

Remedial Strategy

Incoll Management Pty Ltd

On behalf of Rail Corporation NSW

Former Macdonaldtown Gasworks Burren St Erskineville, NSW

> July 2011 JBS 40913 – 15505 Revision 8 JBS Environmental Pty Ltd



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List of Abbreviations

A list of the common abbreviations used throughout this report is provided below.

- As Arsenic
- B(a)P
 Benzo (a) pyrene
- BTEX Benzene, Toluene, Ethylbenzene and Xylenes
- Cd Cadmium
- Cr Chromium
- Cu Copper
- DECCW NSW Department of Environment, Climate Change and Water
- DoP NSW Department of Planning
- DP Deposited Plan
- DQO Data Quality Objectives
- EMP Environmental Management Plan
- EPA New South Wales Environment Protection Authority
- Hg Mercury
- HIL Health Based Investigation Level
- LOR Limit of Reporting
- MAH Monocyclic Aromatic Hydrocarbon
- Ni Nickel
- NoW NSW Office of Water
- OCP Organochlorine Pesticide
- PAH Polycyclic Aromatic Hydrocarbons
- Pb Lead
- PIL Phytotoxicity Based Investigation Level
- PCB Polychlorinated Biphenyls
- PQL Practical Quantitation Limit
- QA/QC Quality Assurance/Quality Control
- RPD Relative Percentage Difference
- RWVP Remediation Validation Work Plan
- SAR Site Audit Report
- SAS Site Audit Statement
- TPH Total Petroleum Hydrocarbons (C₆-C₉ and C₁₀-C₃₆)
- Zn Zinc



Executive Summary

Introduction

JBS Environmental Pty Ltd was engaged by Incoll Management Pty Ltd (Incoll) on behalf of the Rail Corporation NSW Environmental Projects Unit (RailCorp) to prepare a Remedial Strategy for the proposed remediation of the former Macdonaldtown gasworks, located at Burren Street, Erskineville, NSW. The site is identified as part Lot 50 in Deposited Plan 1001467 and occupies an approximate area of 7750 m².

Background

Several contamination investigations have been completed at, or near, the site which have identified soil and groundwater contamination caused by historic site activities. In August 2000 the Site was declared by the NSW Environment Protection Authority (EPA) to pose a Significant Risk of Harm (SRoH) to human health and the environment. Remediation works will be required on the site to remove and /or manage the source of contamination and to render the site suitable for ongoing industrial land use.

A Remedial Action Plan (RAP) was prepared for the site and documented in '*Remedial Action Plan, Former Macdonaldtown Gasworks, Burren Street, Erskineville*, (CH2M Hill 2007). A Site Audit Report (SAR) was subsequently prepared on the RAP by ENSR Aecom and documented in '*Site Audit Report on Remedial Action Plan, Former Macdonaldtown Gasworks Site, Burren Street, Erskineville, NSW*, (ENSR Aecom 2008) which concluded that, in the opinion of the appointed Site Auditor, "...the remediation approach presented in the RAP could be implemented ...in order for the site to be made suitable for the future use for rail-related activities".

An Environmental Assessment (EA) is being prepared for submission to gain approval for the remedial works under Part 3A (Major Infrastructure and Other Projects) of the *Environmental Planning and Assessment Act 1979.* Consideration as part of the EA process, of the full range of project impacts (*e.g.* on the heritage significant items present on site, and to residents in the surrounding area), has determined that the range of remediation technologies specified in the RAP may not be ideally suited to the project, despite being technically feasible.

Additionally, since completion of the RAP and SAR, RailCorp has advised that a section of land contained within the Chullora Railway Workshops, located on Worth Street, Chullora, NSW is available for treatment of soil excavated from the site.

Objectives

Further to the RAP prepared for the site, the objectives of this Remedial Strategy document are to:

- refine the consideration of available/suitable remediation methodologies, based on additional information obtained since the completion of the RAP and SAR;
- outline RailCorp's requirements for each identified likely applicable methodology;
- facilitate a thorough assessment of available remedial options; and
- provide additional site and contaminant data to commence detailed remediation planning.



Consideration of RAP and Additional Remedial Options

A re-assessment was undertaken of the preferred remedial methods provided in the RAP and consideration was also given to alternate remedial methods that could be applied to the site. The re-assessment was primarily based on newly available information on the likely project constraints and requirements, commissioned as part of the EA process.

As part of the assessment of possible options, *in-situ* chemical oxidation and thermal treatment, listed in the preferred remedial methods in the RAP, were determined to not be appropriate for the project. *Ex-situ* remediation of material by thermal treatment, while being technically suitable, was ultimately assessed to be cost prohibitive given the anticipated volume requiring treatment. Remediation of impacted areas by *in-situ* chemical oxidation was originally considered given the lesser need for widescale site disturbance, but was ultimately considered to be poorly suited to the tight clay and shale subsurface present, and also poorly suited to remediation of free tar impacted source zones as identified on the site.

Based on the range and distribution of contamination present, the assessment concluded that no single remedial method provided a solution that was cost effective, timely and appropriate to the site as a whole. Rather, based on the characteristics of the material encountered, the assessment identified four methods that could be used in combination on the site. The four applicable methods comprising the remedial strategy are summarised in **Table 1** below, along with the corresponding suitable materials.

Remediation Method	Likely Suitable Materials		
Excavation and off-site disposal of untreated material	Most cost effective on material unlikely to achieve validation criteria through treatment in a timely manner AND classed as 'Restricted Solid Waste' or lower for off-site disposal		
Excavation and treatment of material for on-site reuse	Most cost effective on material likely to achieve validation criteria through treatment in a timely manner. Onsite treatment method = bioremediation		
Excavation and treatment of material for off-site disposal	Most cost effective on material unlikely to achieve validation criteria through treatment in a timely manner AND classed as 'Restricted Solid Waste' or higher for off-site disposal. Material may be treated on site or off site. Onsite treatment method = bioremediation, Off-site treatment method = cement stabilisation		
In-situ capping of impacted material	Only acceptable where excavations have reached their practicable extent		

The two treatment methods considered most appropriate for the project are bioremediation and cement stabilisation. Based on the anticipated quantities of material and indicative program of works, the configuration of treatment works will involve bioremediation of material on the Macdonaldtown site and cement stabilisation works on the off-site treatment area.

Pre Remediation Documents and Requirements

Implementation of the remedial strategy will also require endorsement of the site specific leachability criteria documented in '*Derivation of Site Specific Leachability Criteria – Former Macdonaldtown gasworks, Burren Street, Erskineville, NSW*' JBS Environmental Pty Ltd, by the appointed Site Auditor.

Additionally, prior to the commencement of remedial works, the following documents will require completion:



- Structural Engineers assessment of retaining structures required around Southern Gasholder;
- Geotechnical specification for treated material to be reused on site;
- Dilapidation Studies on adjacent structures as required;
- A Remedial Health and Safety Management Plan (RHSMP); and
- A Remedial Works Validation Plan (RWVP).



1 Introduction

1.1 Introduction

JBS Environmental Pty Ltd (JBS) was engaged by Incoll Management Pty Ltd (Incoll) on behalf of the Rail Corporation NSW Environmental Projects Unit (RailCorp) to prepare a Remedial Strategy, as prescribed in this document, for the proposed remediation of the former Macdonaldtown Gasworks, located at Burren St, Erskineville, NSW (the site).

Several contamination investigations have been completed at or near the site, which have identified soil and groundwater contamination caused by historic site activities. In August 2000 the Site was declared by the NSW Environment Protection Authority (EPA) to pose a Significant Risk of Harm (SRoH) to human health and the environment. Remediation works will be required on the site to remove and /or manage the source of contamination and to render the site suitable for ongoing industrial land use.

A Remedial Action Plan (RAP) was prepared for the site and documented in '*Remedial Action Plan, Former Macdonaldtown Gasworks, Burren Street, Erskineville*, (CH2M Hill, 2007). A Site Audit Report (SAR) was subsequently prepared on the RAP by ENSR Aecom and documented in '*Site Audit Report on Remedial Action Plan, Former Macdonaldtown Gasworks Site, Burren Street, Erskineville, NSW*, (ENSR Aecom, 2008) which concluded that, in the opinion of the appointed Site Auditor, "...the remediation approach presented in *the RAP could be implemented ...in order for the site to be made suitable for the future use for rail-related activities*".

An Environmental Assessment (EA) has been prepared for the remedial works under Part 3A (Major Infrastructure and Other Projects) of the *Environmental Planning and Assessment Act 1979.* Consideration as part of the EA process of the full range of project impacts (*e.g.* on the heritage significant items present on site, and to residents in the surrounding area), has determined that the range of remediation technologies specified in the RAP may not be ideally suited to the project, despite being technically feasible.

Additionally, since completion of the RAP and SAR, RailCorp has advised that a section of land contained within the Chullora Railway Workshops, located on Worth Street, Chullora, NSW is available for treatment of soil excavated from the site.

1.2 Objectives

In light of these additional project considerations, this Remedial Strategy document has been prepared to:

- refine the consideration of available/suitable remediation methodologies, based on additional information obtained since the completion of the RAP and SAR;
- outline RailCorp's requirements for each identified likely applicable methodology;
- facilitate a thorough assessment of available remedial options; and
- provide additional site and contamination data to commence detailed remediation planning.

This Remedial Strategy document provides a brief summary of the characterisation of site contamination as presented in previous investigations completed on the site. Additionally, this document contains the results of additional field investigations conducted by JBS to facilitate planning for remediation of the site, including a water treatment trial; pump tests



on shallow wells; assessment of clay content in soils; analysis of samples for leachable concentrations of contaminants; and a cement stabilisation trial.

1.3 Structure of the Document

This document has been prepared as an addendum to the RAP (CH2M Hill, 2007) prepared for the site. It should be read in conjunction with the RAP. For ease of use and to avoid unnecessary repetition, where no change is proposed to the strategy, reference is made to the relevant section in the original RAP. The purpose of this document is only to provide further information on the options for remediation of the site, and is not intended to address all the requirements of a RAP as specified in *'Contaminated Sites: Guidelines for Consultants Reporting on Contaminated Sites'* (NSW EPA, 1997).

This document is structured as follows:

- Section 2 provides a brief summary of site details and the contamination status;
- Section 3 details RailCorp's objectives for the remediation and required environmental performance and details the geotechnical, heritage and other major constraints of the project;
- Section 4 summarises the remediation approach recommended in the Remedial Action Plan prepared for the site, assessment of other credible options and a revised remedial strategy incorporating these additional options;
- Section 5 summarises the Remediation Acceptance Criteria to be adopted as part of the revised remediation strategy and the validation sampling plan where deviations from the RAP specified program may be acceptable;
- Section 6 details works required prior to the commencement of remediation;
- Section 7 summarises the pre-treatment requirements for remedial works;
- Section 8 details the consideration for management of perched groundwater in fill and use of a Water Treatment System during the remediation works;
- Section 9 details considerations for treatment of soil;
- Section 10 details considerations of off-site disposal of untreated and treated soil; and
- Section 11 details the anticipated remedial timeframe.



2 Summary of Site Information

On the basis of the complete environmental data set available for the site, a site conceptual model has been prepared and is summarised in the following sections. Full details on site description, history and previous results are provided in previous reports.

Figures showing site location, area and sampling locations are provided as **Figures 1** to **5**.

2.1 Geology

Review of the Sydney Geological Series Sheet 9130 (C. Herbert, 1999) indicates that the geological formation underlying the Site is the Wianamatta Group Ashfield Shale comprising black to dark-grey shale and laminite.

Previous investigations on the site as reported in CH2M Hill (2007b) have identified three general soil types on the site including fill materials, natural soils and tar impacted fill and natural soils. Each is described in more detail following.

2.1.1 Fill Material

Based on the findings of previous investigations as reported in CH2M Hill (2007b), the fill materials identified at the Site have been grouped as follows:

- <u>Ash and Coke Gravels</u> observed across the majority of the Site in surface and near surface layers from ground level to approximately 0.5 m depth;
- <u>Reworked Clays</u> observed in subsurface layers in some site areas between 0.5m depth to approximately 1.5 m depth. This material was observed in the majority of areas as general filling;
- <u>Sands and Gravels</u> observed in subsurface layers in some site areas between 0.5m depth to approximately 1.5 m depth. This material was observed in the North East, South Central and Gas Purifier areas;
- <u>Gravelly Sand and Clay with Minor Ash</u> observed in surface and subsurface layers in some site areas from ground level to approximately 3.5 m depth. This material was predominantly observed in the South West area of the Site as general filling; and
- <u>Gravel, Sand and Demolition Wastes</u> observed in the fill embankment of the Retaining Wall and inside the annulus of the Northern Gasholder. This material was observed to mainly consist of sandy gravels and some ash gravels. It also consisted of demolition wastes and rubble including bricks, metal pipes, tiles, fibrous cement sheeting and Asbestos Containing Material (ACM) and other building rubble in a gravely sand matrix.

2.1.2 Natural Soil

Based on the findings of previous investigations as reported in CH2M Hill (2007b), the natural soil materials identified at the Site have been grouped as follows:

- <u>Silty Clay</u> observed generally from between 1.5 m depth to approximately 2.5 m depth. This material exists across the majority of Site areas. This horizon was predominantly a saturated zone, which sustained the perched groundwater system;
- <u>Red/Grey Mottled Clay</u> observed generally from between 2.5 m depth to approximately 4.0-6.0 m depth. The soil profile is consistent with a Red



Podzolic soil, being moderately to highly plastic, stiff to very stiff, moist and mottled red/grey; and

• <u>Weathered Shale</u> – observed underlying the natural clay. This material grades from extremely weathered to moderately weathered at depths of up to 10 m depth. At depths beyond 6 m, fracturing of the material is common.

2.1.3 Tar Impacted Fill Material and Natural Soils

A number of areas of fill/natural soil materials were observed to be impacted by tar and were summarised in CH2M Hill (2007b). The tar impacts have been categorised as follows:

- <u>Soil/fill impacted by free tar</u> consisting of soil and fill materials impacted to a high degree with black ooze, highly odorous, liquor type material;
- <u>Tarry soils</u> consisting of soil and fill materials with minor tar impacts and moderate odours; and
- <u>Dark Stained Impacts</u> this material was observed as dark brown to black staining in the deep soils and Weathered Shale within the soil pores and shale fractures zones underneath the Southern Gasholder. This material was also moderately odorous.

Soil/fill impacted by free tar was reported by CH2M Hill (2007b) to be predominately associated with former gasworks infrastructure, which include the:

- <u>Tar Wells</u> shallow subsurface and deep natural soils immediately adjacent to these two structures;
- <u>Northern Gasholder</u> deep natural soils immediately adjacent to the brick base annulus; and
- <u>Old gasworks pipework</u> inside pipes and immediately adjacent fill/natural soils.

Tarry soils are present in localised areas, and given free tar have not migrated significant distances from gasworks infrastructure, there appears to be spatial separation between former gasworks infrastructure and tarry soils. Notably there is a layer of highly impacted soils (free tar impacts) surrounding these structures followed by less impacted tarry soils. Tarry soils are located in the following areas:

- <u>Tar Wells</u>, Northern Gasholder and Gas Purifier soil and fill surrounding these source areas in surface/subsurface fill and deeper natural soils;
- <u>Retort</u> fill and deep soil across the majority of this area;
- Gas Purifier Sandy fill and deeper soils; and
- Localised impacted fill observed in one localised pocket in the Northeast Area.

Dark stained impacts were also reported in CH2M Hill (2007b) to be associated with deep soils below the base of the annulus of the southern gasholder. The dark stained impacts were considered likely to be secondary sources within the strata in localised areas associated with the Southern Gasholder.

2.2 Hydrogeology

The groundwater system in the proximity of the site was reported in CH2M Hill (2007b) as existing as a shallow perched groundwater layer and a deep bedrock layer. The shallow groundwater exists within fill materials and silty clay above the natural clay (as shallow as



1m below ground surface), and the deeper groundwater exists within the Ashfield Shale bedrock under semi-confinement.

The groundwater flow direction was reported in CH2M Hill (2007b) to be toward the south/southeast for both shallow and deep groundwater systems. However, it was considered that flows were likely to be influenced by underground structures, including the gasholders annuli and underground waste pits and services associated with gasworks sites. It is possible there may be some interconnectivity between the shallow and deep groundwater systems given the similar direction of flow gradient.

Flow velocities within the shallow groundwater were estimated in CH2M Hill (2007b) to be 6.2-13.7 m/year, while within the deep groundwater are 12.2-36.5 m/year. However, these values do not correlate with the lateral extent of the plume, given that gasworks operations began over 100 years ago. With respect to estimated flow velocities for shallow groundwater it is noted that this layer is likely to be a local layer only and, based on review of test pit logs, flow characteristics are likely controlled by layers of high permeability (*i.e.* gravel, sand or poorly compacted materials) interdispersed between predominantly clay fill.

Based on the results of laboratory analysis of groundwater samples collected from on site and off site wells, the SKM (2006) report stated that:

"The shallow plume appears to begin near the northern boundary of the Former Cleaning Shed and Gasworks areas and extend in a south-west direction of some 75m. The data indicate that the down-gradient edge of the plume is located at the East Hills Line at the southern edge of the site boundary. The lateral extent of the plume appears to be confined in the west to the sewer main located adjacent to the rear boundary of the residential properties, while to [sic] the plume is estimated to extend 50m to the east of the former tank area.

The extent of the middle to heavy-end hydrocarbon plume in the deeper aquifer appears to be larger than the shallow aquifer. While the northern, eastern and western boundaries of the plume are similar to the shallow plume, the down-gradient extent of the plume appears to cover a distance of some 160m from the former tar tank area, with its edge near the southern boundary of railway land along Railway Parade. The data indicate that the deep aquifer plume is located entirely on railway owned [sic] land."

Based on the assumption that the gasworks operations commenced more than 100 years ago, the RAP (CH2M Hill, 2007) notes that the flow velocity values provided in SKM (2006) do not correlate with the measured lateral extent of the plume (reported in the above quotation to extend 75m downgradient and 50m laterally in shallow groundwater, and 160 m downgradient in deep groundwater). It is also noted that the SKM (2006) report acknowledges the same point that, based on the measured extent of the shallow and deep groundwater plumes migrating from the site, the permeability rates estimated from slug tests appear to be an order of magnitude greater than actual rates as suggested by the dimensions of the plume. No explanation was provided for the difference in these permeability rates.



3 Remediation Objectives and Requirements

3.1 Overview

The RAP (CH2M Hill 2007) states that RailCorp wish to remediate the Site such that the following long term objectives can be met, including:

- Removal of the health risks to future site users;
- Removal of the risks to environmental receptors; and
- Allow the beneficial use of the Site for rail related activities.

3.2 Regulations and Approvals

Approval for the remedial works are being sought under Part 3A (Major Infrastructure and Other Project) of the *Environmental Planning and Assessment Act 1979*. Adherence must be made to any requirements placed on the works as part of that Approval.

3.3 RAP and SAR

The remedial works must be undertaken in accordance with the requirements of the RAP (CH2M Hill, 2007) and SAR (ENSR Aecom, 2008) prepared for the site.

3.4 Specific RailCorp Requirements

3.4.1 Requirements Near Site Boundaries

RailCorp has indicated the following requirements with regard to excavations near site boundaries:

- The recommendations for structural retentions along the western site boundary, to be specified by the RailCorp appointed Structural Engineer, must be implemented. Use of batters along this boundary may restrict the removal of impacted material, and is considered inadequate at this end of the site given its proximity to both the former Northern Gasholder source area and the neighbouring Burren Street residences; and
- Temporary batters, constructed in accordance with the recommendation provide in the Geotechnical Report, may be used along the northern, eastern and southern boundaries, provided it will not restrict the progress of excavations or adversely impact structures on adjacent properties.

3.4.2 Operational Environmental and Safety Requirements

Any remediation approach undertaken at the site must satisfy the requirements of RailCorp's environmental and safety policies.

RailCorp's Environmental Policy requires commitment to:

- Minimising use of natural resources;
- Adhering to the principles of ecologically sustainable development;
- Complying with applicable environmental legislation and regulations;
- Effectively managing environmental impacts; and
- Implementing environmental risk management at operational and strategic levels to minimise environmental impacts.

RailCorp's Safety Policy requires commitment to:

• Provide employees, contractors and labour-hire employees a safe environment from injury and workplace-related illnesses; and



• Manage risks proactively to minimise incidents.

Remediation methods applicable to the site are discussed in further detail in **Section 5**, and **Table 5.2** includes the environmental performance requirements of each remediation option to comply with RailCorp's environmental and safety policies.

3.4.3 Closure Requirements

Macdonaldtown

At the completion of remediation works RailCorp's long term objectives for the site should be met. The site should be in a condition such that a Site Audit Statement (SAS) can be prepared declaring the site suitable for commercial/industrial land use. Any Environmental Management Plan (EMP) required to achieve the long term objectives, and preparation of the SAS, should limit the commitments placed on RailCorp in terms of the scope of ongoing works and restrictions on future use. RailCorp must be consulted prior to finalising the requirements of any EMP for the site.

At the completion of remedial works, a network of monitoring wells is also required on the site, sufficient to enable the completion of a program of Monitored Natural Attenuation (MNA) on the site. MNA will be undertaken in accordance with Section 10.3 of the RAP (CH2M Hill 2007) or an Auditor endorsed Groundwater Management Plan for the site.

<u>Chullora</u>

A baseline assessment of the treatment area to be used within the Chullora Railway Workshops must be completed prior to commencement comprising investigation of the potential chemicals of concern (PCOC) in soil and groundwater in accordance with relevant guidelines. The scope of the groundwater investigation as part of the baseline assessment should be appropriate to the nature of the proposed treatment works and their likelihood to result in contamination reaching the water table. At the completion of works, the treatment area must be restored to its pre-treatment works condition. A post treatment assessment of the area, of the same scope as the baseline assessment, will therefore be required to demonstrate that no adverse impacts have occurred or to determine the scale of restoration works.

3.5 Heritage Requirements

Archaeological assessment was undertaken on the site by City Plan Heritage and documented in the report '*Macdonaldtown Gasworks Archaeological Test Excavation Report for RailCorp*' August 2010 Ch10-009. The recommendations of the report pertinent to the remediation works include:

- machine excavation around the State heritage listed Southern Gasholder, is to be adequately planned and supervised to avoid any damage to the structure;
- in the Northern Gasholder providing the bricks forming the annulus are not contaminated, where possible, the bricks should remain *in-situ*. Should removal of the bricks be required to remediate contamination, the removal is to be undertaken with care, so that if possible, the bricks can be cleaned and reinstated. Archaeological monitoring to record the removal, the depth of the annulus and its general construction details should be undertaken during excavation and removal of the annulus. Archival recording of the top of the annulus should be carried out prior to any removal with minimal excavation recommended. If the bricks need to be removed and cannot be reused, the northern gasholder should be represented in some similar form;



- Prior to full scale remediation works an archaeological excavation in the area of the Retort House should be carried out for evidence of the retorts. This would entail machine stripping, hand excavation and recording of the uncovered features to Heritage Branch standards; and
- Prior to full scale remediation works, an archaeological excavation in the area probably containing the footprint of the superintendent's residence should be carried out to record the structural layout of the building. This would entail machine stripping, some hand excavation and recording of the uncovered features to Heritage Branch standards.

3.6 Geotechnical Requirements

Geotechnical testing was undertaken on the site by Pells Sullivan Meynink (PSM) and documented in the draft report '*Macdonaldtown Gasworks Remediation Project Geotechnical Investigation*' July 2010 PSM1444.R1. A summary of recommendations relevant to the completion of the bulk of excavation works is provided below, however, the original report should be referred to for full detail on earthworks and reinstatement requirements.

The report noted that excavation of fill on the site can be undertaken using conventional earth moving equipment, while rock hammers may be required for excavation of shale. **Table 3.1** summarises the recommendations made in relation to acceptable temporary batter slope angles.

Distance between crest		Maximum batter height			
and structure	Geotechnical unit	<4 m	<6 m	<8 m	
More than the batter	Fill	1H: 1V	1.5H:1V	2H:1V	
height	Residual clay	1H:1V	1H:1V	1.5H:1V	
	Fill	2H: 1V	2H:1V	2.5H:1V	
Less than the batter height	Residual clay	2H: 1V	2H:1V	2H: 1V	

 Table 3.1: Recommended Temporary Batter Slope Angles

The report also noted that:

"Temporary batters where structures are located within 1.0 m of the crest should be inspected by a suitably qualified geotechnical engineer during excavation.

Staging of construction to limit the plan extent of the excavation may be able to be adopted to result in localised steeper batter slopes. If such steeper slopes are required additional specific advice should be sought."

The remediation works will need to ensure these slopes are maintained at all times. It is noted that the areas of identified contamination may extend below 8 m, and may also extend up to the site boundary. Provision should therefore be made for the installation of retaining structures to support the walls of the excavation in these areas.

The report also states that fill material present on the site, from a geotechnical perspective only, may be developed "...to allow placement of the majority of the excavated and remediated material as engineered fill". Should treated material satisfy contaminant requirements for reuse on site, it will be necessary to demonstrate that the treated material is suitable from a geotechnical perspective for use as engineered fill on land to be used for railway purposes.

The design of the remedial works should also take into consideration any advice provided from the appointed Structural Engineer in relation to requirements for ground stabilisation prior to and during excavation, particularly in the vicinity of the gasholders and underground services on the site.

3.7 Air Emissions Requirements

An assessment of air quality impacts was undertaken by JBS and is documented in the report titled '*Air Quality Assessment, Remediation of Former Macdonaldtown Gasworks*', Revision F, Reference 40913- 15136, dated August 2011 (JBS 2011a).

A summary of recommendations relevant to the completion of the bulk of excavation works on the Macdonaldtown site is provided below in **Table 3.2**, however, the original report should be referred to for full detail on requirements of dust and odour suppression and monitoring:

Table 3.2	Summary of Required Air Quality Controls				
Site Area / Activity	Proposed Air Quality Control				
Surface soil	Reduction of exposed <i>in-situ</i> materials to 400m ²				
Excavations	Dust suppression by hourly watering of all surfaces				
Retaining Wall fill	Reduction of exposed <i>in-situ</i> materials to 25m ²				
materials	Dust suppression by hourly watering of all surfaces				
Former gasworks area	Enclosure of excavations works, minimum required extent of enclosure shown on Figure 7				
	Treatment of enclosure emissions prior to discharge.				
Soil treatment -	Enclosure of treatment works				
bioremediation	Treatment of enclosure emissions prior to discharge				
Haulage Road use	Dust suppression by hourly watering of all surfaces				
Groundwater	Relocation of water treatment plant to central section of the site as far removed from the Burren Street site frontage as possible, recommended location shown on Figure 7 Enclosure of any areas used for splash filling of water treatment plant Ventilation from water treatment plant to be filtered Prevention of groundwater accumulating within excavations on the site. This may be achieved by pumping water out of the excavations as it infiltrates or if possible by pumping groundwater from adjacent wells				

An assessment of air quality impacts on the alternate treatment site was also undertaken by JBS and is documented in the report titled '*Air Quality Assessment, Remediation of Former Macdonaldtown Gasworks – Chullora Material Receipt Facility*', Revision F, Reference 40913- 15137, dated June August (JBS 2011b).

The recommendations provided in JBS (2011b) relevant to the completion of soil treatment works, were that:

- All soil treatment works on the site be undertaken within temporary enclosure, including the storage of soils over the curing period, and that it be operated under negative pressure conditions to enable extraction and treatment of air emissions from the enclosure; and
- "The extent of exposed coal tar impacted soils is to be minimised to a surface area of 150m². Other coal tar impacted soils stockpiled / bioremediated on the site are to be covered to prevent odour emissions".

The original report should be referred to for full detail on requirements of dust and odour suppression and monitoring.



4 Remedial Strategy Refinement

4.1 Extent of Soil Remediation and Strategy Proposed In RAP

The anticipated extent of remediation as specified in CH2M Hill (2007) is reproduced in **Table 4.1** following. Active remediation was considered by CH2M Hill to be only required for the free tar and impacted fill and soils on site. Requirements for groundwater remediation were discussed in Section 5.6 of the RAP. Reference should be made to **Figures 3** and **4** for the location of each nominated remediation area.

The proposed remediation is a source removal approach, with the intention that the excavation of source material in each area be completed to the extent practicable. Where heritage or geotechnical constraints are encountered such that the practicable limit is unable to remove the full extent of source material, then an *in-situ* management strategy may be implemented in these areas subject to endorsement by RailCorp and the Site Auditor. Areas where such constraints may be encountered include soil at depth in the vicinity of both the western site boundary and the former northern gasholder. Any strategy to contain source material in-situ will need to be compliant with the requirements of *'Guidelines for the Assessment of On-site Containment of Contaminated Soil*' (Australian and New Zealand Environment and Conservation Council, 1999).

The extent of remediation proposed was considered sufficient to protect the health of the future site users by removing or controlling the identified unacceptable health risks.

The contaminants that were considered to drive the health risks were the known carcinogens including benzene and B(a)P. These contaminants were considered to have a direct relationship to the tar source material and the ash/coke surface fill. Therefore, remediation of the tar sources and the ash/coke fill was proposed to mitigate the health risks these contaminants pose to the receptors. This is intended to mitigate the risks by reducing the potential for direct exposure.

Remediation of the tar sources (*e.g.* those accumulated in the Northern Gasholder annulus and the Tar Wells) and remediation of tarry impacted soils (*e.g.* from within the Retort and Gas Purifier areas) was also considered necessary to protect the environmental values of the site groundwater by a reduction in contaminant mass.

Remediation Area	Impacted Area	Estimated Volume (m ³)	Description	RAP Preferred Remedial Approach
Tar wells	Base annulus and immediate area	1000	Soil / fill impacted by free tar	Stabilisation for off-site disposal
	Tar well contents	100	Tar sludge	Off-site disposal as liquid waste with pre-treatment as required to improve handling
Northern gasholder	Base annulus and immediate area	2100	Soil / fill impacted by free tar	Stabilisation for off-site- disposal to landfill, with pre- treatment as required to improve handling
	Gasholder contents	640	Impacted water	Off-site disposal as liquid waste
		320	Tar sludge	Off-site disposal as liquid waste with pre-treatment as required to improve

Table 4.1: Summary of Remedial Extent and Strategy (as adapted from Table 5.1, CH2M Hi	ill,
2007)	



Remediation Area	Impacted Area	Estimated Volume (m³)	Description	RAP Preferred Remedial Approach
				handling
	Buried wastes inside annulus	1900	Demolition materials	Off-site disposal as' Asbestos/Industrial' ¹ (<i>i.e.</i> with pre-treatment to remove free tar or oversize component)
Former gasworks area	Shallow fill / soils	9225	Tarry soils – fill and natural clays	Stabilisation for off-site- disposal to landfill, with pre- treatment as required to improve handling
	Deeper soils	2375	Tarry soils – natural clays and weathered shales	Stabilisation for off-site- disposal to landfill, with pre- treatment as required to improve handling
	TP16 Hotspot	115	Tarry soils – fill and natural clays	Stabilisation for off-site- disposal to landfill, with pre- treatment as required to improve handling
Site surfaces	Site surfaces		Ash and coke gravels	Off-site disposal to landfill
Retaining wall		1765	Gravel sand and demolition wastes	Off-site disposal to landfill. With processing (<i>i.e.</i> segregating oversized component) recycling or beneficial reuse may also be an option
Hotspots	BH14	100	Fill and natural clays	Off-site disposal to landfill
	MW13s	140	Fill	Offsite disposal to landfill
	MW04s	100	Fill and natural clays	Off-site disposal to landfill
Pipework	Varying across site	Unknown	Tar / scrap metal	Treatment to remove tar from pipework. Tar to be disposed to landfill, scrap metal possibly disposed to landfill as demolition waste

4.2 Additional Remediation Options Consideration

Further to several detailed studies on the implementation of the RAP, it was decided that based on the range and volume of materials requiring remediation on the site, the proposed strategy should incorporate alternate options for remediating the site to those provided in the RAP. JBS was requested to review the potential for the following additional treatment/management options to be incorporated into the remedial approach for the site:

- Excavation of soil for ex-situ treatment by bioremediation for on site reuse; and
- *In-situ* capping of impacted materials.

An assessment of the technical and overall suitability was undertaken for the additional options listed above, and is summarised in **Table 4.2**.

¹ Asbestos Contaminated Special Waste or Restricted Solid Waste under DECC 2008



Each of the possible remedial options has been assessed for each of the three contamination issues requiring assessment on the site, namely:

- Free tar present in disused infrastructure, in fill, soils and shale underlying the site;
- Ash and tar materials considered to be acting as source material for unacceptable levels of groundwater contamination; and
- Asbestos impact potentially contained throughout fill materials on the site

Table 4.2 also includes assessment of two of the four remedial methods preferred in theRAP (CH2M Hill 2007), specifically:

- excavation and off-site disposal of soil without treatment; and
- excavation and off-site stabilisation of soil for off site disposal.

No further consideration was made of chemical oxidation of material and thermal treatment for off site disposal, despite their inclusion as preferred methods in the RAP (CH2M Hill, 2007). Chemical oxidation was considered unsuited to the particular contaminants of concern, and to the low permeability of the soil and rock units underlying the site. Thermal desorption was considered technically feasible, but less suitable, given the proximity of sensitive land uses to the site (and potential off site treatment location), and likely costs for the low volume of material to be treated.

4.3 Revised Remedial Strategy

Based on the additional data and review of other likely remedial methodologies, the remedial strategy for the site is presented in the following flowchart (**Flowchart 4.1**) and summarised in **Table 4.3**.

Remediation Method	Likely Suitable Materials
Excavation and off-site disposal of untreated material	Most cost effective on material unlikely to achieve validation criteria through treatment in a timely manner AND classed as 'Restricted Solid Waste' or lower for off-site disposal
Excavation and treatment of material for on-site reuse	Most cost effective on material likely to achieve validation criteria through treatment in a timely manner. Onsite treatment method = bioremediation
Excavation and treatment of material for off-site disposal	Most cost effective on material unlikely to achieve validation criteria through treatment in a timely manner AND classed as 'Restricted Solid Waste' or higher for off-site disposal. Material may be treated on site or off site. Onsite treatment method = bioremediation, Off-site treatment method = cement stabilisation
In-situ capping of impacted material	Only acceptable where excavations have reached their practicable extent

Table 4.3: Summary	of Remedial Strategy and Suitable Materials	
Tuble Hor Summar	of Remeater Strategy and Suitable Platerials	

Further consideration each element of the Remedial Strategy is provided in Table A.



Table 4.2: Evaluation of Soil Treatment and Management Options

	in or som meatment and management options			
Consideration	Option 1 – Excavation and off site disposal of untreated material	Option 2 – Excavation and on-site treatment for on site reuse (Bioremediation)	Option 3 – Excavation and off-site treatment for off site disposal (stabilisation)	Optio
Use of permanent solutions &	Permanent solution: Yes	Permanent solution: Yes	Permanent solution: Yes	Perma
alternative technologies or	Alternate technology: No	Alternate technology: No	Alternate technology: No	Alterna
resource recovery technologies	Resource recovery technology: No	Resource recovery technology: Yes	Resource recovery technology: No	Resour
Satisfy DECC (2006) preference for treatment as a principle element	No	Yes	Yes – although may be undertaken as off site treatment at Chullora ²	No
Suitable materials	Only materials that are classed as Restricted Solid Waste, or lower, for off site disposal.	Only materials that are likely to be treated by bioremediation and likely to achieve validation criteria within both a reasonable time frame and cost. Unlikely to be suited to material impacted by free tar or asbestos	Only materials that are classed as Restricted Solid Waste, or higher, for off site disposal. Unlikely to be cost effective on materials classed as General Solid Waste for off site disposal	Only a reache excava
Ability to achieve validation	Validation able to be achieved with certainty	This option has the potential to achieve validation	This option has the potential to achieve validation	Validat to ong
Ability for treated material to be reused on site	-	This option has the potential to produce material suitable for reuse however, heavily impacted materials may encounter treatment difficulties and potential failure of validation criteria could occur	This option has the potential to produce material suitable for reuse however, the relatively high concentrations and nature of hydrocarbons (heavy end TPH / PAHs) suggests treatment difficulties and potential failure of validation criteria relating to leachability could occur	-
OH&S considerations	OH&S issues during remediation able to be reasonably managed. Intensive odour control will be required for nearby site users	OH&S issues during remediation able to be reasonably managed. Intensive odour control will be required for nearby site users	OH&S issues during remediation able to be reasonably managed, intensive odour control will be required for nearby site users	OH&S
Timing & staging requirements	No significant time delays	Uncertain – generally extended timeframes required for bioremediation.	Uncertain – generally extended timeframes required for bioremediation. Dependent also on capacity of stabilisation system, and available area as will require 7-10 days for curing of each treatment batch	No sig factor remed
Geotechnical requirements	Water table reduction in fill required and stabilisation	on of Southern Gasholder and any other structures to be	retained required (common to all options)	
	Upfront costs: low	Upfront costs: moderate	Upfront costs: low	Upfron
	Water Treatment Costs (common to all potential op	ptions: uncertain will also depend on costs, if any, for dis	posal of groundwater generated during dewatering)	
Cost effectiveness	Long Term Costs: High, given large areas of the site where material would be classed as Hazardous Waste, Restricted Solid Waste or Special Waste for off-site disposal	Long Term Costs: Uncertain – will be dependent on treatment timeframes	Long Term Costs: Moderate	Long T on-goi associa incider
	Total Costs: Comparatively High	Total Costs: Uncertain, but likely to be less than disposal of treated material to landfill	Total Costs: Uncertain , but likely to be less than disposal of untreated material to landfill	Total C Presen the lar
	Disposal site (<i>i.e.</i> Landfill) will require appropriate	If treated material to be disposed to off site: disposal site (<i>i.e.</i> Landfill) will require Immobilisation Approval from DECCW, disposal site will also need approval to receive waste	Treated material to be disposed to off site: disposal site (<i>i.e.</i> Landfill) will require appropriate EPL ¹ for waste disposal application to land	The co manag will ne the RA
Compliance with applicable or relevant appropriate requirements (ARARs)	EPL ¹ for waste disposal application to land for that class of material	If material to be treated at Chullora an EPL ¹ for treatment of contaminated soil will be required for Chullora	If material to be treated at Chullora EPL ¹ for treatment of contaminated soil required for Chullora	suitabl of cap Site Au materi
	Water Generated During Dewatering – Trade Waste	e Agreement with Sydney Water if not suitable for reinjed	ction (common to all methods)	
				Requir

ion 4 – In-situ Capping of Impacted soils

manent solution: Yes ernate technology: No ource recovery technology: No

y acceptable where excavations have been ched the practicable limits i.e. where remedial avation cannot removal full extent of source

dation able to be achieved with certainty, subject ngoing management.

&S issues able to be reasonably controlled

significant delays , however program will need to for in time for construction of capping before nediation is complete

ront costs: moderate

g Term Costs: low, however party responsible for going management will maintain the liability ociated with human health and environmental dents linked to breach of the containment area

al Costs: comparatively low

sence of containment area may reduce value of land in the future

e consent authority may need to accept the *in-situ* nagement of contaminated material and a method need to be nominated for legal enforcement of RAP. There will need to be a responsible party able to be nominated for on-going management capping and / or containment cell, approval from a Auditor required for strategy to determine which terials suitable to remain on site.

uires on-going management, may limit future mercial use of the site and responsibility if



Consideration	Option 1 – Excavation and off site disposal of untreated material	Option 2 – Excavation and on-site treatment for on site reuse (Bioremediation)	Option 3 – Excavation and off-site treatment for off site disposal (stabilisation)	Optio
Remediation				breach
Protection of Human Health	Overall Good	Overall Good	Overall Average – may not be capable of remediating concentrations present in workable timeframe.	Overal mass, limit fu
			Good - reduction in contaminant mass on site	
			Poor - generation of green house gases during remediation	
	Good - reduction in contaminant mass on site	Good – reduces the leachable concentration of contaminants	Poor - if used without air emission controls then potential for odour generation during remediation	Good - enviro
Environment & heritage	Poor - consumes limited landfill resources	If off-site disposal required as a result of failed treatment, poor - option consumes limited landfill resources	Poor - need to segregate asbestos impacted material from treatment process or incorporate mitigation measures in operational procedures to prevent release /exposure to asbestos fibres during and post treatment	Poor- r therefo manag
			Disposed to landfill, poor - consumes limited landfill resources	
Reputation / community	Consultation required but assume preferable as only certified clean materials will be used to reinstate site	Consultation required but assume preferable as reduces number of large vehicle movements to and from site	Consultation required but assume preferable as only certified clean materials will be used to reinstate site	Consul disrup
Conclusion	Suited only to material impacted by low levels of contamination. Primary limiting factors include: - No licensed facilities in NSW exist to receive material classed as 'Hazardous Waste' for off site disposal - High cost option - High impact on limited landfill resources Not a complete solution	Suited only to material considered suited to achieve validation criteria by bioremediation in a reasonable timeframe. Primary limiting factors include: - bioremediation unsuited for remediation of inorganic and semi-volatile contaminants - Potential cost of off site disposal if validation criteria cannot be achieved after treatment Not a complete solution	May be used as the complete solution, however costs likely to be prohibitive if used on material capable of being reused on site Cost of undertaking treatment by stabilisation are likely to restrict the suitability of this option only to material considered unsuitable for treatment by bioremediation and/or in untreated form classed as Restricted Solid Waste, or higher, for offsite disposal	Accep reachd The pi remov Not a

Notes:

1 EPL: Environment Protection Licence issued by DECCW under the Protection of the Environment Operations Act (1997)

ion 4 – In-situ Capping of Impacted soils

aches occur

rall **Good** but as no reduction in contaminant as, method requires on-going management, may t future commercial use of the site.

d – isolated contaminated material from ironment

r- no reduction in contaminant mass on site refore maintenance of isolation involves ongoing nagement and liability

sultation required but assume preferable as less uption to surrounding area

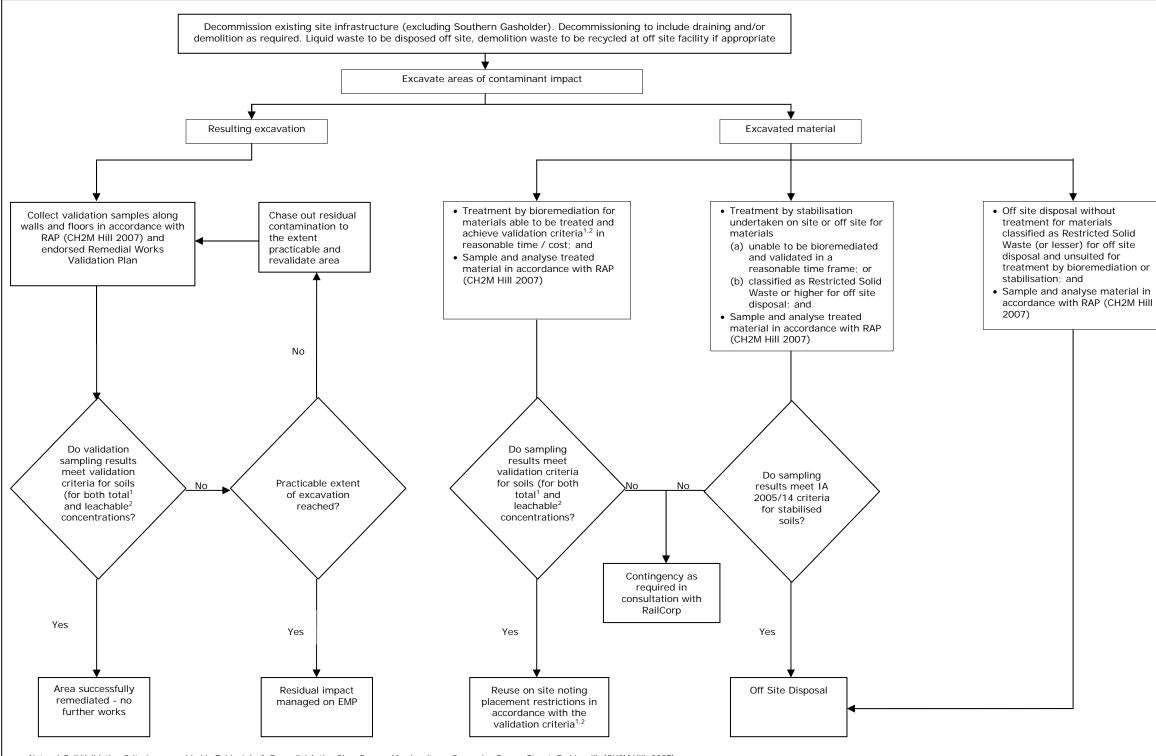
eptable only where the excavations have ched practicable limits

e primary limiting factor is the that source noval is not achieved adopting this option

a complete solution



Flowchart 4.1 Remedial Strategy



Notes: 1 Soil Validation Criteria as provided in Table 4.1 of 'Remedial Action Plan, Former Macdonaltown Gasworks, Burren Street, Erskineville'(CH2M Hill, 2007) 2 Site Specific Leachability Criteria as provided in Table 10 of 'Derivation of Site Specific Leachability Criteria. Former Macdonaldtown Gasworks' (JBS 2010a)



5 Remediation Acceptance Criteria and Validation Plan

Validation data are to be collected to verify the effectiveness of the remediation works and document the condition of the site as being suitable for the proposed future use(s).

Given the nature and extent of remediation works, validation data shall verify that:

- the identified contaminated soils (listed in **Table 5.3** of this document and Section 5 of the RAP) were effectively remediated; and
- any soils / fill materials retained on the site (including any materials excavated and treated for reuse) are suitable for on-going industrial land use.

5.1 Remediation Acceptance Criteria

The RAP (CH2M Hill, 2007) produced for the site included site specific validation criteria for total concentration in soil. The calculated values are depth dependent, based on risks to future site users associated with exposure to vapours and present for four distinct depth ranges:

- Surface (0.0m) to 1.5m below ground surface of finished site;
- 1.5 to 2.5m below ground surface of finished site;
- 2.5 to 4.0m below ground surface of finished site; and
- 4.0 to 8.0m below ground surface of finished site.

Soil validation criteria are summarised in **Table 5.1** following.

Table 5.1: Soil Validation Criteria

Analyte	Depth Range ⁶			
	0-1.5m	1.5-2.5m	2.5-4.0m	4.0-8.0m
Heavy metals	•	•	•	•
As (total)	500	-	-	-
Cd	100	-	-	-
Cr	500 ¹	-	-	-
Cu	5,000	-	-	-
Hg (inorganic)	75	-	-	-
Ni	3,000	-	-	-
Zn	35,000	-	-	-
Monocyclic Aromatic Hydro	carbons			
Benzene	1 ³	1 ³	1 ³	1 ³
Toluene	1.4 ²	2.6	4	7.9
Ethylbenzene	3.1 ²	11.1	17.6	34.8
Total xylene	14 ^{2, 3}	14 ³	14 ³	14 ³
Polycyclic Aromatic Hydroc	arbons (PAHs)	·		
PAHs (total)	100	-	-	-
Benzo(a)pyrene	5	-4	-4	-4
Naphthalene	-	3.8	6.0	11.8
Acenaphthene	-	-4	-4	_4
Fluorene	-	_4	-4	-4
Pyrene	-	_4	-4	-4
Benzo(b)fluoranthene	-	-4	-4	-4
Chrysene	-	-4	-4	-4
Other Constituents	-			
Total Phenol	42,500	-	-	-
Cyanide (complex	2,500	-	-	-
Asbestos	No detection	of fibres in surfac	e soils (0.5m dep	th). No visible

Note: 1. Value is for Cr(VI) and used as a conservative concentration as a preliminary screening value for chromium.



Criteria for toluene, ethylbenzene and xylenes at 0-1.5m are ecological health based.
 Risk based values are lower than laboratory analytical limits of reporting (LORs) and health investigation levels (HILs), therefore less conservative HILs applied to all depths.

4. Not of concern. Based on the outcomes of vapour fate and transport modelling reported in CH2M Hill (2007b) the contaminant was considered to have a low vapour potential at the nominated soil temperature of 15°C.

5. Adopted criteria in CH2M Hill (2007b) on the basis of Australian Contaminated land Consultants Association (2002) 'Asbestos in Soils – Code of Practice'

6. Depth ranges provided in metres below ground level in RAP (CH2M Hill 2007), these will need to be converted to levels relative the required finished ground surface.

Given that the revised strategy allows for material to be reused onsite, site specific leachability criteria were derived in the JBS letter '*Derivation of Site Specific Leachability Criteria – Former Macdonaldtown gasworks, Burren Street, Erskineville, NSW*' Reference JBS 40913-15501, provided as **Appendix B.**

The site specific leachability criteria were derived based on the relevant criteria for protection of groundwater resources in the area, but also incorporating a dilution attenuation factor (DAF) to account for dilution of contaminated groundwater that occurs at the receiving water body, as it becomes mixed with groundwater discharged from the remainder of the catchment. A DAF value of 16.6 was determined for the site, and the resulting site specific leachability criteria for the main contaminants of concern are provided in **Table 5.2** below. The complete list of site specific leachability criteria is presented in **Appendix B**.

Contaminant	Limit of Reporting	Site Specific Criteria for assessment for leachable concentrations in soil ¹
HEAVY METALS		
Arsenic (III/V)	0.1	38.2 / 74.7
Cadmium	0.1	11.6
Chromium (III)		166
Chromium (VI)	0.1	73
Copper	0.1	21.6
Lead	0.1	73
Manganese	1	1328
Mercury	0.05	1.76
Nickel	1	1162
Zinc	1	249
POLYCYCLIC AROMATIC H	IYDRCAOCARBONS	
Benzo(a)Pyrene	0.1	1.7
Naphthalene	0.1	1162
Phenanthrene	0.1	10.3
Anthracene	0.1	0.2
Fluoranthene	0.1	1.7

Table 5.2: Site Specific Leachability RAC (all units in µg/L)

¹ Adopted criteria – 16.6x ANZECC/ARMCANZ 2000 trigger values, 16.6 x the LOR was adopted where ANZECC/ARMCANZ 2000 trigger values are unlikely to be achieved readily by the laboratories

The values presented in **Table 5.2** were derived to be protective of downgradient receptors in the event that contamination leaches out of subsurface soils, infiltrates the water table and migrates off site (**Appendix B**). They are considered to be the upper limits of allowable leachabilities on site. The actual quality of shallow groundwater on the site (**Table 8.2**), is much better than the upper limits set for leachability. Given that the remediation strategy prescribed will ultimately remove the bulk of contaminant mass from the subsurface it is therefore considered unlikely that



the leachability of material in validated areas of the site, or material to be reinstated after treatment will reach the upper limits provided in **Table 5.2**.

5.2 Validation of Soil Remediation Works

The proposed soil validation sampling and analytical program for the revised remedial strategy is outlined in **Table 5.3**. Validation requirements for material to be imported onto site have been included for completeness, noting that the specified testing requirements will need to be undertaken on the material at its source location and prior to arrival on site.

Item	Sampling Frequency		Analytes	
	Excavation floors	Excavation Walls (from each distinct horizon / material type)	Sampling Density	
Remediation Excavation (consistent with RAP)	1 sample per 8.5 m square grid centres	1 sample per 10 lineal metres / 10 m – at depths of between 0 and 1.5m, 1.5 and 2.5m, 2.5 and 4.0m, 4.0 and 8.0m and every 2m below 8.0m depth	N/A	8 metals BTEX PAHs Phenols Cyanide Asbestos
Treated materials prior to off-site disposal or on-site reuse (ALTERNATE TO RAP)	N/A	N/A	7 samples per treatment batch up to 1000 m ³ + 1 additional sample per 200m ³ (or part thereof) in excess of 1000m ³	8 metals BTEX PAHs Phenols BTEX (gw leach) ¹ PAH (gw leach) ¹ Heavy metals (gw leach) ¹
Impacted material to be retained in- situ under newly installed site capping	Confirmatory sampling on 8.5 m square grid centres. Samples analysed at 1.0 m intervals to a depth of 1 m below observed depth of impact		8 metals BTEX (total and GW leach) PAHs (total and GW leach) Phenols Cyanide Asbestos	
Imported material for reinstatement ² : - VENM, ENM soil or material generated and certified under an approved DECCW resource recovery exemption and approved for use by DECCW ²	N/A	N/A	Minimum 10 samples per source site to enable calculation of 95%UCL _{avg} concentrations	8 metals TPH/BTEX PAHs OCPs/PCBs (for VENM only) Asbestos Foreign material (ENM only) pH (ENM only) VCH (ENM only)
(ALTERNATE TO RAP)			For recovered mater be in accordance wit of the relevant exem	ials testing should h the requirements

Table 5.3: Soil Validation Sampling Program

Notes: 1 leachability testing shall be undertaken on treated materials using site specific groundwater to simulate likely potential risks to groundwater.

2 All imported material must be tested and validated at the source location prior to receipt at the Macdonaldtown Gasworks site.

3 any material generated under an approved DECCW resource recovery exemption to be imported to site must be demonstrated as complying with all the requirements of that exemption and must be approved for use in reinstating the site in writing by DECCW.

It is noted that TPH is not included in the analytical suite specified for validation samples, despite the inclusion of petroleum hydrocarbons in the '*principal chemicals of interest at gasworks sites*' in DEC (2005) '*Information for the assessment of former gasworks sites*'. Imray and Langley (Enhealth, 2001)² mention the difficulties in the risk assessment of mixtures such as TPH and refer to two approaches used in a hybrid framework for assessing TPH by The Total Petroleum Hydrocarbon Criteria Working Group (TPHCWG,

² Health-based Soil Investigation Levels, Imray, P. and Langley, A., enHealth (enHealth, 2001).



1997). The first approach investigates the presence of indicator chemicals which are carcinogenic substances such as benzene and PAHs, which if detected require assessment using contaminant specific health investigation levels. The second stage involves assessment against criteria for well defined TPH fractions, which have been derived for $>C_{16}-C_{35}$ aromatics, $>C_{16}-C_{35}$ aliphatics and $>C_{35}$ aliphatics in soil. The analytical suite for the validation program is based on the first approach, i.e. assessing the specific compounds likely to present an unacceptable risk, rather than the TPH mixture itself. This is considered appropriate for the site as the extensive historical dataset indicates that elevated TPH concentrations were primarily comprise of either PAH or BTEX compounds. Conversely TPH concentrations were generally below the detection limit in samples where PAH or BTEX concentrations were low.

Sampling Rationale

The sampling and analytical regime presented in **Table 5.3** is based on the following rationale:

- The wall sampling frequency is based on what is considered to be an adequately conservative lineal frequency capable of detecting residual contamination and being representative of residual materials;
- Treated material sampling frequencies for on site reuse or off-site disposal are designed to ensure reliable 95%UCL_{avg} concentrations are derived; and
- Imported material sampling frequencies exceed the minimum 1 composite sample requirement outlined in EPA 1995, and is based on having sufficient data to generate reliable 95%UCL_{avg} concentrations. Procedure B (EPA 1995) will also be used to confirm that an appropriate number of samples have been obtained from each source type / material type to enable comparison against the appropriate criteria.

Sampling Methodology and Data Quality

Should remain consistent with the requirements of the Section 9 of RAP (CH2M Hill, 2007)

5.3 Application of RAC

The following rules apply to use of RAC in material to be retained on site:

- For impacted material to be retained *in-situ* due to heritage or geotechnical restrictions on the extent of excavation – soils in the area to be retained on site must be sampled and analysed at the frequency specified in Section 5.3, which includes both total and leachable concentrations. Leachable concentrations in all samples must comply with the RAC provided in Table 5.2 due to the potential for contaminant migration. Consultation should be undertaken with RailCorp/Site Auditor on a case by case basis to determine whether these materials are also required to comply with the RAC for total concentrations at specific locations. Where samples fail to meet these criteria a contingency strategy may be required subject to RailCorp/Site Auditor endorsement prior to implementation. Contingency options may include management by capping and ongoing monitoring;
- For material used to reinstate the site following treatment by bioremediationeach treated batch is to be sampled and analysed at the frequency specified in Section 5.3. Leachable concentrations in all samples must comply with the RAC provided in Table 5.2, AND total concentrations in all samples must be compliant with RAC (specified in Table 5.1, as adapted from the RAP, CH2M Hill, 2007).



Where samples fail to meet these criteria a contingency strategy may be required subject to RailCorp/Site Auditor endorsement prior to implementation. Contingency options may include extending the period of remediation, cement stabilisation or disposal to landfill without further treatment; and

3. For material to be treated by cement stabilisation – as this material will require waste classification for off site disposal, each treated batch is to be sampled and analysed in accordance with the requirements of IA 2005/14. Leachable concentrations should be assessed against Specific Contaminant Concentration (SCC) values provided in Waste Classification Guidelines (DECC 2009), AND compliance with the additional criteria listed in **Table 5.4** must be demonstrated. Where samples fail to meet these criteria a contingency strategy may be required and will require RailCorp/Site Auditor endorsement prior to implementation. Contingency options may include retreating of soils to achieve compressive strength or disposal to landfill;

Property	Requirement (IA 2005/14, NSW EPA) ¹
Maximum allowable	PAHs – 13 000 mg.kg ⁻¹ B(a)P – 500 mg.kg ⁻¹ Non-halogenated phenols – 2000 mg.kg ⁻¹ Total Cyanide 4000 mg.kg ⁻¹
Reagent used for stabilisation	Calcium or Magnesium Oxide
Maximum allowable stabilisation ratio	2:1 (<i>i.e.</i> by mass, 2 parts reagent to 1 part untreated soil)
Unconfined compressive strength (UCS)	1 MPa as the 95% lower confidence limit value (as assessed by AS1012.9-1999)

Table 5.4: Soil Criteria (units as specified)

Notes: 1. General Approval of the Immobilisation of Contaminants in Waste' (NSW EPA 2005), Approval Number 2005/14 - Coal Tar Waste from former Gasworks Sites.

5.4 Groundwater Management

No active groundwater remediation is proposed in the RAP (CH2M Hill, 2007). The Monitored Natural Attenuation (MNA) approach is proposed to monitor the concentrations of residual compounds in groundwater following the completion of soil.

It will, therefore, be necessary as part of the soil remediation works to install new groundwater monitoring well on the site in accordance with Section 10.3.3 of the RAP (CH2M Hill, 2007), or any future RailCorp and Auditor endorsed Groundwater Management Plan produced for the site.

Additionally any perched groundwater drained out of the fill material and shallow soil to enable excavation of the areas to be remediated will require treatment prior to off site disposal. Further discussion of the treatment of perched water generated during excavation works is provided in **Section 7**.



6 Pre-Remedial Documentation

A number of pre-remedial studies, plans and approvals are likely to be required for implementation of the RAP (CH2M Hill 2007) and this Remedial Strategies document. These are summarised following.

6.1 Pre-Remedial Studies

The following pre-remedial studies should be completed prior to the detailed consideration and design of remediation works:

6.1.1 Geotechnical and Dewatering Assessments

As the areas requiring remediation extend up to the site boundaries, geotechnical structural retention will be required to secure properties bordering the site. As a minimum it is considered geotechnical controls will be required along the western boundary and in the vicinity of the southern gas holder, but consideration should also be given to the need for retention along the southern and northern site boundaries.

Geotechnical assessment may also be required on treated material proposed for reuse on the site and for disposal to landfill.

Where not included in the existing geotechnical report³ then detailed geotechnical assessments will be required to provide geotechnical parameters for retaining structure design.

Structural engineer design of shoring will be required based on the geotechnical parameters.

Each of the studies shall be undertaken by qualified geotechnical and structural engineers.

6.2 Pre Remediation Works Plans

6.2.1 Remediation Health and Safety Management Plan (RHSMP)

Several potential health and safety hazards are anticipated to be present during the course of the remediation works. These relate to physical hazard posed by the completion of a large scale demolition and earthworks project and chemical hazards associated with the contaminated soils and groundwater underlying the site. A Remediation Health and safety Plan (RHSMP) will be required prior to the commencement of works. As a minimum the RHSMP will need to detail the following items:

- Assessment of potential hazards posed by works including detailed descriptions of potential toxicological impacts from contaminants present in soil and groundwater underlying the site;
- Stipulation of measures to remove hazards (where possible);
- Procedures / controls to be put in place to control hazards where elimination is not possible;
- Any requirement for personal protective equipment to be worn by the site workforce;
- Specific consideration of the PPE and/or operational requirements for maintenance of acceptable working conditions within the proposed enclosure;



- Requirements for pre-works training of the site workforce (*i.e.* Occupational Health and Safety General Induction for all personnel, HAZWOPER training for staff potentially directly exposed to contaminants etc);
- Requirements for occupational monitoring to be completed during the remediation works; and
- Evacuation plans and directions for medical assistance / first aid.

Additional requirements for the RHSMP may be identified during the course of remediation.

6.2.2 Dilapidation Studies

There is a potential for proposed remediation works to have structural impacts on adjoining properties. Pre- and post-works dilapidation studies will be required of adjoining properties to assess for dilapidation as caused by the remediation works.

6.2.3 Remediation Works and Validation Plan (RWVP)

The requirements of this document and the RAP may be achieved by a number of methods by the successful remediation contractor. A Remediation Works and Validation Plan (RWVP) will be required prior to the commencement of works and will require endorsement by the appointed Site Auditor. As a minimum the RWVP will need to detail the following items:

- Compliance with the requirements of the RAP, the Environmental Management Plan (EMP) prepared for the site and this Remedial Strategy document will be met;
- Compliance with any additional requirements arising out of the Part 3A approval;
- The validation program (frequencies and analytical suites) to be adopted for material to remain *in-situ*;
- The validation program (frequencies and analytical suites) to be adopted for treated material to be reused on site;
- The preferred off-site disposal location(s) for material to be disposed to landfill;
- Details of the treatment methods to be adopted, both on-site and off-site;
- An indicative layout of the likely treatment operation relative to site boundaries and adjacent land uses, including the proposed infrastructure and drainage plans on both sites, and on the Chullora site waste reception and handling areas;
- Details of the baseline sampling program to be adopted on the treatment site; and
- Details of the post-treatment sampling program and the criteria to be met at the completion of off-site treatment works.

6.3 Pre-Remediation Approvals

A review of the likely scope of remediation works has been completed. The approvals identified as most likely being required include:

• Development consent as an integrated development under the *Environmental Planning and Assessment Act 1979* from City of Sydney Council prior to commencement of any demolition or excavated works associated with the remediation of the site. The works will be classified as a Category 1 remediation works under SEPP 55 'Remediation of Land';



- Should source removal during remediation require dewatering of the major water bearing zone underlying the site, then a Construction Dewatering Permit under the *Water Act 1912* will be required;
- Approval to discharge treated groundwater generated as part of the remediation works. For discharge to sewer, a Trade Waste Agreement must be entered into with Sydney Water. For discharge into stormwater approval will be required from City of Sydney Council. Approval may also be required for the reuse of treated groundwater on the site;
- An Environment Protection Licence from NSW EPA under the *Protection of the Environment Operations Act 1997* may be required where untreated soil is to be transferred off site for treatment;
- Should the stabilisation method adopted deviate from the specification provided in IA 2005/14, then a Specific Immobilisation approval will be required from the NSW DECCW as per NSW DECC (2008). It is likely that correspondence from the proposed landfill disposal location will be required confirming that the premises are licensed to receive immobilised material;
- If it is intended that material treated at Chullora is to be returned to site as treated material, then licensing by NSW EPA under the *Protection of the Environment Operations Act 1997* will be required;
- Where a substantial quantity of materials require removal from the site generating significant vehicle movements then approval may be required under SEPP11 'Traffic Generating Developments'; and
- WorkCover NSW must be provided with notification seven days prior to any demolition works, or asbestos removal works.



7 Site Establishment and Configuration of Operations

Limits on the area available for remediation on the Macdonaldtown site and available for treatment works on the Chullora site are likely to determine the configuration of works undertaken.

7.1 Macdonaldtown

Based on the remedial strategy summarised in **Flowchart 4.1**, a summary of the proposed scope of works on the Macdonaldtown site is presented in **Table 7.1**. **Figures 6** to **9**, nominate the anticipated remediation areas/depths and likely locations for the associated infrastructure.

Stage	Task	Comments
Preliminary	Project planning and licensing	-
Site Establishment I	Setup of site offices, sediment and erosion controls	-
Remediation Stage I	 1A – assessment/soil sampling of northern boundary retaining wall 1B- construction of internal turning circle, vegetation removal 	Blue area forming the northern boundary of the site as shown on Figure 5 (does not include fill material within the former Northern gasholder)
	1C – excavation/validation of the top 0.5m of fill material of the entire site surface. Transfer of excavated soil to Chullora for treatment prior to disposal to landfill	Yellow areas on Figure 5 (ash & coke gravel fill) do not have a malodorous potential. These soils will be excavated by standard excavation practice to typical depth of 0.5m.
		Consideration may be given to supplementary sampling through this layer prior to excavation, for comparison to remediation acceptance criteria as defined in the RAP (CH2M Hill 2007)
	1D – excavation/validation of hotspots to depths of 1-2m	Hot-spot 'A' at north-east portion of the site (Figure 5) in proximity of former sample location BH14; Benzene impact.
		Hot-spot 'B' at eastern portion of the site (Figure 5) in proximity of former sample location TP16; Free tar impact.
		Hot-spot 'C' at eastern portion of the site (Figure 5) in proximity of former sample location MW04 (Figure 3); Benzene impact.
		Hot-spot 'D' at south-western portion of the site (Figure 5) in proximity of former sample location MW13; Ash and coke impacts
		Excavation/validation of any other hotpot locations identified during the preceding stages
Site Establishment II	Installation of temporary enclosure, associated air extraction/treatment system and water treatment system	Proposed locations shown on Figure 7.
Remediation Stage II	2A- commission air and water treatment system	-
	2B – excavate/validate areas within enclosure. Transfer of excavated soil to Chullora for treatment prior to disposal to landfill	Pink and orange areas on Figure 5 as present within the boundaries of the enclosure
	2C – reinstate enclosure excavation with imported material certified as suitable for the proposed land use and compliant with the	

Table 7.1: Summary of Remediation Works



Stage	Task	Comments
	relevant legislation	
Remediation Stage	3A – excavate/validate areas external to the enclosure. Excavated material unsuited for onsite bioremediation within enclosure to be transferred to Chullora for treatment prior to disposal to landfill	Orange areas on Figure 5 as present external to the boundaries of the enclosure During completion of Task 3A the water treatment may require location based on the extent of chase out excavation required.
	3B –Material assessed as suitable for remediation by bioremediation to be stockpiled for treatment within enclosure	Stockpiles of material awaiting bioremediation within the enclosure will be placed in the areas designated for soil stockpiling as shown on Figure 5 and be maintained in accordance with the EMP for the works (JBS 2011c)
	3C – reinstatement of site using imported material certified as suitable for the proposed land use and compliant with the relevant legislation	-
Disestablishment	Decommissioning of air and water treatment plants, disestablishment of enclosure and site offices	-

A program of controls for odours, gas and dust emissions from the Site and routine monitoring has been designed to mitigate the impact of the proposed works on the surrounding community and environment. Full details of the required management controls and monitoring program are provided in *'Environmental Management Plan, Demolition and Remediation, Former Macdonaldtown Gasworks, Burren Street, Erskineville, NSW*, (JBS 2011c).

The major control to be adopted is for excavation of primary source zones to be completed within an enclosed area. Use of an enclosed area was a recommendation of the Air Quality Impact Assessment (JBS 2011a) required for the control of odour and gas emissions from heavily impacted areas. **Figure 7** shows the likely position of the enclosure. The enclosure shall be a purpose built metal clad or fabric structure sufficiently sized to allow the internal operation of tippers, excavators and associated equipment as required for the stockpiling and handling of soils and any associated equipment. Openings in the enclosure shall be minimised so as to reduce potential for uncontrolled releases. A temporary purpose-built enclosure has been nominated rather than a more permanent engineered constructed building given the:

- need for remediation to occur in a large purpose built structure which can optimise use of space without limiting remediation works or requiring substantial preconstruction design and site preparation or post-remediation demolition and waste;
- need to minimise disruption to the normal operations of the adjacent Macdonaldtown Stabling Yards; and
- post-remediation concept plan for the site is for open space land absent of any new buildings.

It is anticipated that a purpose built temporary enclosure operated under negative pressure conditions that is demonstrated effective through routine monitoring (in accordance with JBS 2011c) will provide control equal to that of an engineered building.

The recommendations provided in '*Air Quality Assessment, Remediation of Former Macdonaldtown Gasworks'*, Revision F, Reference 40913- 15136, (JBS 2011a) require that the temporary enclosure be maintained under a constant negative pressure during working



hours. To this end it will be necessary for the enclosure to be fitted with an emissions control system that will allow for controlled extraction and treatment of air.

The emissions control system will comprise two main elements:

- The collection system which must be capable of maintaining adequate ventilation rates throughout the enclosure under negative pressure conditions. With the exception of fugitive emission that may occur when the enclosure entry points are momentarily opened and closed, the collection system must be capable of extracting emissions out of the enclosure through one central and controlled point; and
- The treatment system external to the enclosure and must be capable of receiving and treating air extracted from within the enclosure. The treatment system should utilise granular activated carbon (GAC) filter(s) with appropriate sizing to remove potentially malodorous or harmful constituents. Figure 7 shows the likely location of the air treatment system.

Any bioremediation proposed as part of the remediation program is to be conducted within the temporary enclosure. Spreading, turning and stockpiling of soil undergoing bioremediation within the zone of influence of the emissions control system will mitigate the potential for offensive odours to migrate beyond the boundaries of the Site. Construction of windrows or biopiles of soil for bioremediation will need to be appropriately sized as per the dimensions of the enclosure. It is proposed that bioremediation occurs after the completion of excavation works within the enclosure, therefore consideration should be given to changes in the air treatment system that may be required to accommodate likely increases in emission and particulate loading rates. In the event that increased loading rates are expected during bioremediation, re-commissioning of the air treatment system will be required. Design of the emissions control system should ensure an efficiency capable of meeting OH&S requirements for air quality within the enclosure, and/or specify requirement for personal protective equipment (PPE) requirements for the exposed workforce. Any requirements for PPE within the tent will need to be included in the RHSMP, as detailed in **Section 6.2.1**.

Based on the available dataset it is not anticipated that free tar will be encountered external to the temporary enclosure. However as a precaution, a contingency has been included in JBS (2011c) for such an occurrence and requires that the following is completed prior to the disturbance of free tar impacts external to the enclosure:

- Works in the area cease until the Remediation Consultant has determined the appropriate controls for that location, including OH& considerations for the remediation workforce;
- The free tar material remains securely covered for the duration between the first encounter with the material and the remedial excavation works. Plastic sheeting, soil, steel plates or other appropriate cover may be used for this purpose;
- Wherever possible any sampling required to characterise the free tar and adjacent material should be undertaken prior to the commencement of remedial excavations for the free tar. Appropriate disposal locations should also be confirmed prior to commencement; and
- Where free tar is encountered external to the temporary enclosure the material will need to be excavated and transported to a licensed landfill for treatment or disposal.



JBS (2011c) notes that the controls required are likely to be specific to the occurrence of the free tar, however lists the following as possible contingency actions:

- Temporary windscreens installed around the perimeter of the free tar area prior to remediation, in conjunction with wetting of the exposed surfaces during remediation; or
- Delineation of the area, relocation (and operation) of the enclosure over each occurrence, as required, once all remediation works inside the original footprint have been completed to the extent practicable.

Additionally these contingencies may also require execution in those areas within the enclosure footprint where the excavation cannot be safely extended to the point of successful validation without impacting the stability of the enclosure. This includes where the excavation begins to impinge upon allowable batter slopes, as summarised in **Section 3.6**, or past the line of temporary shoring.

Any additional infrastructure on the treatment site required by the contractor to meet the conditions of the RAP (CH2M Hill 2007), the REMP (JBS 2011c) and this Remedial Strategy document will need to be documented in the RWP to be prepared as per **Section 6.2.3**.

7.2 Chullora

A summary of the proposed scope of works on the Chullora site is presented in **Table 7.2** including likely plant required. **Figure 10** nominates the anticipated site setup.

Stage	Task	Comments
Preliminary	Project planning and licensing	-
Site Establishment	Conduct baseline environmental assessment of treatment site	-
	Setup of site offices, sediment and erosion controls	-
	Installation of temporary enclosure and associated air extraction/treatment system. Installation of cement stabilisation plant within the enclosure	Proposed locations shown on Figure 10.
Treatment	A- commission air and water treatment system B – receive materials for treatment. Onsite stockpiling until minimum treatment volume achieved - Proposed areas for stockpiling of on Figure 10 and soils in this ar- maintained in accordance with th for the works (JBS 2011c)	
	C – once minimum volume achieved treatment/validation of soils by cement stabilisation within the enclosure	Treated soil is to remain within the enclosure for the duration of the curing period and until validation results confirm successful stabilisation has occurred.
	D- stockpiling of treated/validated soil external to enclosure until removal off-site is possible	Any treated material removed from site to be transferred directly to a an appropriately licenced landfill for disposal. Proposed areas for stockpiling of shown on Figure 10 and soils awaiting transfer in this area to be maintained in accordance with the EMP for the works (JBS 2011c)
Disestablishment	Decommissioning of air treatment plants, disestablishment of enclosure and site offices	-
	Conduct post works environmental assessment of treatment area	-

Table 7.2: Summary of Treatment Works



A program of controls for odours, gas and dust emissions from the Site and routine monitoring has been designed to mitigate the impact of the proposed works on the surrounding community and environment. Full details of the required management controls and monitoring program are provided in the EMP (JBS 2011c).

The major control to be adopted is for the treatment of impacted soil to be undertaken within an enclosed area, including storage of treated soil over the curing period. Use of an enclosed area was a recommendation of the Air Quality Impact Assessment (JBS 2011b) required for the control of odour and gas emissions from the cement stabilisation process. **Figure 10** shows the likely position of the enclosure. The enclosure shall be a purpose-built metal clad or fabric structure sufficiently sized to allow the internal operation of tippers, excavators and associated equipment as required for the stockpiling and handling of soils and any associated equipment. Openings in the enclosure shall be minimised so as to reduce potential for uncontrolled releases. A temporary purpose-built enclosure has been nominated rather than a more permanent engineered constructed building given the:

- need for treatment to occur in a large purpose built structure which can optimise use of space without limiting remediation works or requiring substantial preconstruction design and site preparation or post-remediation demolition and waste;
- need to minimise disruption to the normal operations of the adjacent Chullora Rail Yards; and
- need for the area to be restored to its original condition on completion of the treatment program.

The recommendations provided in (JBS 2011b) require that the temporary enclosure be maintained under a constant negative pressure during working hours. To this end it will be necessary for the enclosure to be fitted with an emissions control system that will allow for controlled extraction and treatment of air. The emissions control system will need to comprise a collection system and treatment system consistent with that required on the Macdonaldtown site as described in **Section 7.1**.

It is anticipated that a purpose built temporary enclosure operated under negative pressure conditions and one that is demonstrated through routine monitoring (in accordance with JBS 2011c) will provide effective control equal to that of an engineered building.

Stage	Description of Works	Major Equipment								
-	Receipt, Stockpiling, Treatment and	Pug Mill – size to be determined								
	Disposal of Soil	20T excavators – external to enclosure								
		20T excavator – internal to enclosure								
		Air treatment system, including:								
		- Diesel generator								
		- Extraction Fan (2 x 1.5m diameter)								
		- Granular activated carbon filter								
		Semi trailers arriving and departing from site								
		Water Truck								
		Tipper trucks								

Table 7.3: Summary of Treatment Works

Any additional infrastructure on the treatment site required by the contractor to meet the conditions of the RAP (CH2M Hill 2007), the REMP (JBS 2011c) and this Remedial Strategy document will need to be documented in the RWP to be prepared as per **Section 6.2.3**.



8 Pre-Treatment Requirements

Table 8.1 provides additional detail on the requirements of the various elements of the revised remedial strategy and identifies the following pre-treatment works potentially required for the remediation of site:

- Pre-treatment of tar sludge for off-site disposal by either heating or inclusion of additives such as fly ash to improve handling;
- Extraction of tar contents from gasworks pipes manually or by other methods such as using heat or chemicals;
- Lowering moisture content, crushing and/or homogenisation of fill, soil and shale impacted by free tar or displaying other tar impacts;
- Segregation of oversized materials in demolition waste and fill obtained from existing retaining walls and in the vicinity of hotspots at BH14, MW13S and MW04S;
- Removal of impacted water in below ground infrastructure on the site (Northern and Southern Gasholders, tar wells etc) and as required in areas to be excavated.

8.1 Tar Wells and Northern Gasholder

Tar sludge present in the tar wells and Northern Gasholder will most likely be removed from site by specialised vacuum trucks licensed to transport liquid waste. Under these circumstances the tar sludge may be treated *in situ* to improve handling and be pumped directly into the vacuum trucks without the need for a separate above ground treatment area. The heating or requirement for additives to enable pump out of tar sludge will be dependent on several site specific conditions at the time of remediation including volume and depth of tar sludge, proportion of soil or other inclusions, size and power of pump utilised and capability of the disposal truck to maintain the handling properties of the sludge during transport. The Contractor will be responsible for ensuring that OH&S, environmental and/or planning controls are adequately addressed for the proposed tasks.

Extraction of tar contents from gasworks pipes may be undertaken as follows:

- Where the pipes require preservation, then treatment in the form of heating or additives to mobilise and extract the contents; or
- Where the pipes need not be preserved, and assuming proper soil and water controls are in place, then it may be possible to sever the pipes into smaller sections for manual extraction of the contents.

The Contractor will be responsible for ensuring that OH&S, environmental and/or planning controls are adequately addressed for the proposed task.

8.2 Free Tar and Impacted Fill

Pre-treatment works on free tar and tar impacted fill, on either site, will need to be undertaken in a contained area to prevent the uncontrolled spread of contamination and release of vapours. As a minimum sediment and erosion controls should be provided to achieve this end. Additionally where the disturbance of tar sludge occurs outside the tented enclosure (as discussed below and in **Section 3.7**), management controls will also be required to prevent unacceptable releases of contaminants in air or odour. This may include capture of emissions and treatment if necessary.

Pre-treatment of free tar and tar impacted fill soil and bedrock may require lowering the moisture content of the material, crushing the excavated material and/or homogenisation



to enable stabilisation. The pre-treatment works are capable of completion using conventional earthmoving equipment such as excavators and backhoe loaders.

The following applies to any pre-treatment works required on materials excavated from the Northern Gasholder and tar wells further north, and all other coal tar impacted areas, (as described in the Air Quality Assessment completed for the works, JBS 2010a):

- Pre-treatment works undertaken on the Macdonaldtown site on free tar and tar impacted soils will need to be enclosed to prevent unacceptable odour impacts beyond the boundary of the site;
- Pre-treatment works undertaken within the boundaries of the designated treatment area within the Chullora Railway workshops, must ensure that no more than 150m² of coal tar impacted soil is uncovered at any one time. The restriction of exposed surface area of soil on the Chullora treatment site is required in order to prevent unacceptable odour impacts on the surrounding area in accordance with the Air Quality Assessment for the Chullora treatment site (JBS 2010b); and
- The clay content of soil successfully remediated through cement stabilisation generally lies between 60 and 80 % by mass. One of the three soil samples tested in the recent benchscale stabilisation trial contained a clay content of 87%, and when treated with 5, 12.5 and 20 % cement, failed to achieve the required compressive strength specified in IA 2005/14 (NSW EPA 2005) for cement stabilisation. Based on the results of the benchscale trial, it is considered that the materials containing a clay content greater than 80% are likely to occur in natural soil layers encountered at depth. Therefore it may be necessary to obtain a specific immobilisation approval from DECCW for cement stabilisation to allow a lower UCS value to be adopted for the process. Alternately some of the material capable of being treated may require homogenisation prior to treatment. Any material used in the homogenisation process to achieve the required composition in the material for treatment must be derived from the Macdonaldtown site.

Segregation of oversized particles will be required on those materials to be treated by cement stabilisation or bioremediation, and is generally completed through mechanical screening. However, consideration should be given to optimising any segregation works to minimise the amount of handling required of the coal tar impacted material prior to treatment. Consideration should be given to:

- The OH&S and PPE requirements for mechanical screening of fill potentially containing asbestos;
- Management of odour impacts;
- Off –site disposal requirements of the segregated oversized particles, which may
 not be suitable for disposal as building and demolition waste in accordance with
 DECC (2008) if impacted by free tar or other contaminants post sorting.
 Segregated oversized materials that as assessed as 'impacted' may require a
 waste classification for off site disposal.
- The attachment of high speed shredders on the screening plant, given that much of the fill and residual soil underlying the site is predominantly clay. The high speed shredder would be faster than a conventional shaker screen under these conditions and would simultaneously cause the break up of clay clumps in the screened material.

Requirements of the pre-treatment of impacted water from within the northern gas holder and tar wells is discussed in **Section 8.**

					Pre-treatment Requirements		Primary Treatment Requirements					
Site Area	Material Type	Extent of Impact	Expected Quantity (m ³)	Remediation Method	Possible Pre-treatment Required	Waste Classification (subsequent to pre- treatment if required)	Primary Treatment Options	Anticipated Waste Classification Subsequent to Treatment / Immobilisation Approval				
	Tar Sludge Contained within Bas Gasholder		320	Liquid Waste Disposal	Potential pre-treatment to improve handling – heating or Hazardous Waste (Liquid) use of additive such as fly ash		-	-				
	Impacted Water	Contained within Gasholder	640	Liquid Waste Disposal, disposal to sewer or on site reuse	Extraction from gasholder	Liquid Waste	Pass through on site water treatment plant	Suitable for disposal to sewer or on site beneficial reuse				
Northern gasholder	Soil / fill impacted by free tar	Base annulus and proximate soils (within pink shaded area on Figure 4 to a depth of 8m-10m)	2100	Treatment by bioremediation treatment for onsite reuse or landfill disposal OR Treatment by stabilisation to apply NSW DEC immobilisation approval (Approval #2005/14)	Potential options to improve handling if required - lowering moisture content, breakdown clay clods to expose higher surface area	Hazardous	Stabilisation or bioremediation treatment– Stabilisation to apply NSW DEC immobilisation approval (Approval #2005/14)	Restricted Solid Waste or General Solid Waste (may also be assessed for suitability for on site reuse)				
	Demolition	Buried inside Gasholder					Segregation of free tar, asbestos	_s Hazardous Waste (free tar)				
	Waste	annulus (blue shaded area on Figure 4)	1900	Landfill disposal or Recycling	-	Special Waste (asbestos)	containing materials and oversize particles	Special Waste (asbestos)				
							· · · · · · · · · · · · · · · · · · ·	General Solid Waste (Non-Putrescible)				
	Shallow Tar Impacted Soil and Fill	Lateral extent of orange shaded area on Figure 4 to a depth of at least 4m	9225		Physical amendment to break	Hazardous or Restricted Solid Waste		Restricted Solid Waste or General Solid Waste (may also be assessed for suitability for on site reuse)				
Retort and Surrounding Former Gasworks Source Areas	Deep Tar Impacted Natural Soil	Lateral extent of pink shaded area on Figure 4 in the vicinity of boreholes BHE and BHF to a depth of 8m-10m	2375	Treatment by bioremediation treatment for onsite reuse or landfill disposal OR Treatment by stabilisation to apply NSW DEC immobilisation approval	down material with high clay content (<i>i.e.</i> lowering moisture content, breakdown clay clods to expose higher surface area) Likely alternative site for pre-	Hazardous or Restricted Solid Waste	Stabilisation or bioremediation treatment Stabilisation to apply NSW DEC immobilisation approval	Restricted Solid Waste or General Solid Waste (may also be assessed for suitability for on site reuse)				
	Tar Impacted Contamination hotspot at TP16 location	Lateral extent of green shaded area on Figure 4 to a depth of 1 m-2 m	115	(Approval #2005/14)	treatment and remedial treatment	Hazardous or Restricted Solid Waste	(Approval #2005/14)	Restricted Solid Waste or General Solid Waste (may also be assessed for suitability for on site reuse)				
Existing Site Surfaces	Ash/ Coke Fill	Lateral extent of shaded area on Figure 4 to a depth of at least 0.5 m	2950	Stabilisation or bioremediation treatment for onsite reuse or landfill disposal. Application of NSW DEC Immobilisation approval (Approval #1999/05)	Potential options to improve handling if required - lowering moisture content, breakdown clay clods to expose higher surface area	Hazardous	Stabilisation or bioremediation treatment Stabilisation to apply NSW DEC immobilisation approval (Approval #2005/14)	Restricted Solid Waste or General Solid Waste (may also be assessed for suitability for on site reuse)				
Retaining Wall	General Fill and demolition waste	Entire Northern boundary (shaded blue on Figure 4)	1765	Landfill disposal, Beneficial Reuse or Recycling	segregation of oversize material	sGeneral Solid Waste	-	-				
Contamination Hotspots	Impacted Fill at locations	Lateral extent shown as green shaded area on Figure 4 to a depth of 1 m-2 m	340	Treatment by bioremediation treatment for onsite reuse or landfill disposal OR Treatment by stabilisation to apply NSW DEC immobilisation approval (Approval #2005/14)	segregation of oversize material	gregation of oversize materials Hazardous (Approval (Approval #2005/14)		Restricted Solid Waste or General Solid Waste				
Site Wide	Old Gasworks Pipes	Varied	Unknown	Separate landfill disposal of empty pipework and tar contents	Chemical or manual extraction o tar contents, steam, or other cleaning of pipe work	fHazardous (tar) and General Soil Waste Non-Putrescible (clean pipe sections)	-	-				
Site Wide	Fill and natural soil materials	NA	Unknown	Beneficial reuse, or as required: Stabilisation or bioremediation treatment for onsite reuse or landfill disposal.	segregation of oversize materials Hazardous		Stabilisation or bioremediation treatment Stabilisation to apply NSW DEC immobilisation approval (Approval #2005/14)	Restricted Solid Waste or General Solid Waste				
Deep excavations proximal to source area	Residual tar sources – subsequent to source removal	Unknown	Unknown	Treatment by bioremediation treatment for onsite reuse or landfill disposal OR Treatment by stabilisation to apply NSW DEC immobilisation approval (Approval #2005/14)	Potential options to improve handling if required - grinding to expose higher surface area	Hazardous	Stabilisation or bioremediation treatment	Restricted Solid Waste or General Solid Waste				

Table 8.1:	Revised Remedial Strategy - Requirements of Pre – Treatment and Primary Treatment
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9 Considerations for the Water Treatment System

Based on observations recorded during previous testing on the site, excavation of fill material as part of the remediation program may encounter significant volumes of perched groundwater. A water treatment system (WTS) is proposed as part of the remediation strategy to process groundwater generated either during excavation, or pumped out of the fill prior to excavation. The objective of using the WTS would be to treat the collected groundwater to an acceptable level, such that it may be discharged to stormwater, to sewer or used in operation *e.g.* as wheel wash water or for dust suppression.

9.1 Rates of water requiring treatment

Test locations encountering significant seepage in testpits are summarised in **Table 9.1** and displayed on **Figure 11.** Flow velocities within the shallow groundwater were estimated in CH2M Hill (2007) to be between 6.2 and 13.7 m/year, determined through slug tests conducted in shallow groundwater wells on the site.

Test Location	Termination Depth (m BGL)	Depth Extent of Fill (m BGL)	Fill description	Depth of Apparent seepage zone (m)
TPA (CH2M Hill Phase I & II ESA 2000)	2.0 (R)	0 - 2.0	 0 - 1.5: FILL, Light brown medium grained sand with some gravel, loose unconsolidated moist. 1.5 - 2.0: FILL, Free tar product migrating out of bricks seams, some clay, plastic, wet, tarry type odour. 	1.5 - 2.0
TPC (CH2M Hill Phase I & II ESA 2000)	1.6 (R)	0 - 1.6	0 - 1.6: FILL, Bricks, minor clay, sand, very wet, visible hydrocarbon sheen, hydrocarbon odour, tar visible on bricks. (BRICK WALL TP EASTERN SIDE)	0 - 1.6
TP03 (CH2M Hill 2006)	4	0 - 2.8	0- 0.3: FILL gravelly Sand, brown, dark brown, dark grey, ash and crushed sandstone, ballast, clinker, ash 0.3 - 1.5: FILL gravelly Clay, firm, medium plasticity, dry, orange and grey, fine to coarse gravel, subround to subangular, ballast, coke. Water ingres at 0.6m, fast 1.5 - 2.8: FILL clayey Silt (original surface?), low plasticity, dark grey, wet	0.6 - 2.8
TP06 (CH2M Hill 2006)	2.7	0 - 2.3	 0 - 0.3: FILL gravelly Sand, ash, clinker and coke, gravel subround to angular, fine to coarse grained 0.3 - 0.5 FILL sandy Clay, light brown to orange, firm, low plasticity, some clinker 0.5 - 1.4: FILL (original surface?) clayey gravel, orange/brown, medium to coarse grained, sandstone, grey-black subangular clinker GROUNDWATER INGRES AT 0.6m 1.4 - 2.3: FILL clayey Silt (original surface?) dark grey/brown, wet 	0.6 - 2.3
TP07 (CH2M Hill 2006)	2.2	0.2 - 1.7	0.2 - 0.5: FILL gravelly Sand, black to dark grey, dry, ash and coke, gravel fine to medium grained, of ash, coke, clinker 0.5 - 0.9: FILL clayey Sand, brown, slightly gravelly, wet, half bricks, gravel fine to coarse grained, subround to subangular, sandstone, shale 0.9 - 1.3: FILL gravelly Clay, grey and red mottles, very firm, subangular to subround sandstone and shale 1.3 - 1.7: FILL clayey Silt (original surface?), soft	0.5 - 0.9

Table 9.1: Summary of Observed Seepage in Fill



TP10 (CH2M Hill 2006)	4.1	0 - 3.2	 0 - 0.8: FILL gravelly Sand, dark brown, dark grey or black, of ash, coke, coal, gravel medium to coarse grained, angular to subangular of same material, presence of bricks, round cobbles 0.8 - 0.95: FILL single layer of red brick 0.95 - 1.25: FILL Sand, dry medium grained, yellow sand, massive concrete boulder at 1.1 m BGS 1.25 - 1.75: FILL gravelly Clay, orange with grey mottles, dry, medium to coarse grained, subangular to subround, of shale (WATER INGRESS at 1.7m) 1.75 - 2.25: FILL sandy Gravel, grey, wet, of shale 2.25 - 3.2: FILL possibly reworked natural material, orange with grey mottles, very firm, gravel medium to coarse grained, angular shale 	1.75 - 2.25
TP11 (CH2M Hill 2006)	4	0 - 3.2	0 - 0.3: FILL gravelly Sand, brown, black, dry, ash and coke, rootlets in top 0.1 m 0.3 - 0.8: clayey Sand, yellow, orange, wet 0.8 - 3.2 FILL clayey Sand, interbeded with grey and red mottles, firm clay at 0.8 - 0.9 mBGS and 1.2 - 1.3 m BGS, reworked natural material	0.3 - 0.8
TP15 (CH2M Hill 2006)	4.1	0 - 2.5	0 - 0.3: FILL gravelly Sand, loose 0.3 - 1.7 FILL ash and gravel water and free tar at 1.1m bgs at brickwork footing 1.7 - 2.5 FILL sandy Clay (original surface?), brown to green, high plasticity	1.0 - 1.7
TP16 (CH2M Hill 2006)	3.8	0 - 1.5	0 - 0.2: FILL gravelly Sand, dark grey, black, dry, ash and coke 0.2 - 1.5: FILL gravelly Sand, orange and light brown, dry, with cobbles, sandstone GROUNDWATER INGRESS at 0.2 m	0.2 - ?

Results of recent pump testing by JBS, detailed in **Appendix A**, determined the hydraulic conductivities at MW37S and MW42S, to be between 5.09 x 10^{-6} m/s and 6.65 x 10^{-6} m/s.

Anticipated inflows along the north boundary of the site based on recent pump test results may be as high as between 300 m³ per day and 400 m³ per day at the commencement of excavation works, based on the following conservative assumptions:

- saturated fill is present between 1 and 6 m depth across the entire site;
- continuous hydraulic connectivity in groundwater present in fill across the site; and
- excavation of fill material would require pumping along the entire northern boundary of the site (approximately 140m in length).

The PSM (2010) Geotechnical Report provides no assessment on the impact of draining the perched water table on settlement on the site and surrounding properties. Based on the water bearing zones listing in **Table 9.1** it appears that the zones containing significant volumes of perched groundwater are restricted to layers of fill. As these layers are likely to be limited to the extent of the site, the volume of inflows are anticipated to reduce over time. In the event that no reduction of inflows occurs, the remediation works should consider the impacts of settlement on the site and surrounding area.

Additionally, surface water is likely to have accumulated in underground infrastructure associated with the former gasworks including the Southern Gasholder and Tar Wells. Estimates of likely volumes of impacted water contained in subsurface structures are estimated as follows:

- Northern Gasholder 640 m³ (CH2M Hill, 2007);
- Southern Gasholder 1875 m³ (based on a diameter of 20 m and assumed depth of 6 m, filled with water); and
- Tar Wells- 50 m³ (conservative assumption allows for half of 100 m³ of tar well contents, reported in CH2M Hill 2007, to be filled with impacted water).



Some allowance should also be made for impacted water contained in disused service trenches, pits and pipeworks that may remain *in-situ*.

9.2 Anticipated Influent Quality and Required Effluent Quality

Table 9.2 summarises the range of contaminant concentrations detected in groundwater sampled from the site. In viewing **Table 9.2** it should be noted that the majority of groundwater generated from remedial works at the site is likely to be derived from the fill layers, where contaminant concentrations were generally less than those detected in samples of groundwater collected from the underlying natural shale.

	Criteria	Shallow Grou	ndwater Con	centrations	Deep Groundwater Concentrations						
Analyte	ANZECC 2000	Range (µg/L)	Location of Maximum	Site Area	Range (µg/L)	Location of Maximum	Site Area				
Electrical Conductivity	200-300 ¹ (μS/cm)	442 - 2010 (μS/cm)	MW35S	Central northern	717 – 3820 (μS/cm)	MW03D	Central Southern				
Cd	0.2	nd - 2.6	MW13s	Southwest	nd - 1.5	MW06d	Gasholders				
Cr(total)	-	nd – 15	MW04s	South Central	nd - 7	MW04d	South Central				
Cu	1.4	nd – 220	MW42s	Northeast	0.001 - 208	MW42d	Northeast				
Pb	3.4	nd – 174	MW42s	Northeast	nd - 140	MW03d	South Central				
Ni	11	nd – 10	MW04s	South Central	nd - 92	MW36d	Offsite				
Zn	8	0.033 - 1,570	MW13s	Southwest	0.015 - 869	MW42d	Northeast				
Cyanide (total)	7	0.02 - 0.479	MW20s		nd - 14.9	MW03d					
Benzene	950	nd – 704	MW07s		nd - 14,000	MW03d	South Central				
Toluene	-	nd – 117	MW07s		nd - 792	MW03d	South Central				
Ethylbenzene	-	nd - 213	MW07s	Gasholders	nd - 317	MW03d					
Total Xylenes	550 (o & p)	nd - 417	MW07s	1	nd - 5,010	MW03d					
Total PAHs	16 (naphthalene)	nd - 1,677 (naphthalene 1,460)	MW07s		nd - 4,208 (naphthalen e 3,840)	MW07d	Gasholders				

 Table 9.2: Summary of Groundwater Contamination

Note: 1 typical range of EC in NSW lowland rivers as provided in ANZECC 2000 Table 3.3.3

9.3 Required Effluent Quality

Table 9.3 summarises the acceptable concentrations for disposal to sewer under an Industrial 'Trade Waste' Agreement with Sydney Water as provided in the Sydney Water Brochure '*Industrial Customers – Acceptance Standards and Charging Rates for 2010-11*'. A copy of this brochure is provided as **Appendix D**. These criteria are likely to be the required performance criteria for any WTS used on site, noting that in providing approval for discharge into sewer Sydney Water may alter allowable concentrations based on the estimated rates and TDS concentrations. All notes associated with **Table 9.3** should also be considered in determining the requirements of the systems performance.

Discussions with City of Sydney (CoS) staff indicate that should it be necessary to discharge treated groundwater to stormwater, then an application must be made to CoS demonstrating that the water to be discharged:

- Is clear (i.e. turbidity of less than 20 NTU);
- Is free of visible suspended sediment (i.e. total suspended solids concentration less than 50 mg/L);
- Has no visible oil or grease film (i.e. oil and grease less than 10 mg/L);
- Has a pH value between 6.5 and 8.5; and



• Has been assessed to not adversely impact the visual character of the receiving water body i.e. concentrations of any other contaminants of concern occur at levels which do not represent a risk to the receiving water body (i.e. Alexandra Canal).

For treated groundwater to be reused on site *e.g.* as wheelwash water or for dust suppression, compliance with the criteria provided in **Table 9.3** should be sufficient. Consideration may also be given to the use of site specific values, or modified site specific values where appropriate. The current values for site specific leachability criteria are provided in the JBS letter '*Derivation of Site Specific Leachability Criteria – Former Macdonaldtown gasworks, Burren Street, Erskineville, NSW*' Reference JBS 40913-15501, provided as **Appendix B.**

A WTS was trialled on the site, with the methods and results documented in the JBS Letter Report '*Groundwater Treatment Trial, Former Macdonaldtown Gasworks*', dated 05 August 2010 Reference 40913 - 15534. A copy of this letter is included as **Appendix C**.

Overall it was noted that samples of the treatment system effluent were reported to contain very low concentrations of all contaminants of concern at the site. The average concentration of all WTS effluent samples is included in **Table 9.3**, and indicates that with the exception of arsenic, the concentrations in treated water samples complies with the relevant 'Trade Waste' acceptance criteria where available, noting, however, that the full suite of analytes listed in **Table 9.3** was beyond the scope of the trial. With respect to arsenic it was considered that used of an acid washed granular activated carbon (GAC) filter could result in arsenic concentrations within acceptable limits.

Notwithstanding this difference, it appears that the WTS used in the treatment trial could be scaled up for use in the remediation process to enable discharge of treated groundwater to sewer. While the specifics of the system to be used during the remediation program will require on site refinement based on the actual groundwater quality and quantity encountered, the details of the successfully implemented trial system are summarised in the following sections.

It is also noted that when applying for the trade waste agreement, justification should be provided for not undertaking analysis for the full suite of analytes listed in **Table 9.3**, rather a reduced testing suite should be recommended comprising the contaminants of concern at the site i.e. PAHs, heavy metals and water quality parameters and others as required based on the site historical groundwater data.

Contaminant	DISPOSAL TO SEWER – Trade Waste Acceptance Criteria ^{1,3,4,18}
Acetaldehyde	5
Acetone	400
Aluminium	100
Ammonia	100 ⁵
Arsenic	1
Barium	5
Biological Oxygen Demand (soluble)	100 ¹⁷
Boron	100
Bromine	5
Cadmium	1
Chlorinated Phenolics	0.05 ⁸
Chlorine	10
Chromium	39
	5
Cobalt	5
Copper	5 1 ¹⁰
Cyanide	
Fluoride	206
Formaldehyde	30
General pesticides (excludes OC and OP)	0.1 ¹¹
Herbicides and defoliants	0.1
Iron	50
Lead	2
Lithium (Specified Systems only)	10 ¹²
Manganese	10
Mercaptans	1
Mercury	0.03
Methyl Ethyl Ketone	100
Molybdenum	100
Nickel	3
Organoarsenic compounds	0.1
рН	7 to 10
Petroleum Hydrocarbons (flammable)	10 ^{20, 13,16}
Benzene	0.17
Toluene	0.5
	1
Ethylbenzene	
Xylene	1 1 ⁸
Phenolic Compounds	
Polynuclear aromatic hydrocarbons	5
Benzo(a)pyrene	-
Naphthalene	-
Phenanthrene	-
Anthracene	-
Fluoranthene	-
Propionaldehyde	5
Selenium (total)	5
Silver	5
Sulphide	5
Sulphite	50
Temperature	38°C
Thiosulphate	300
Tin	10
Total Dissolved Solids	500 ^{14,18}
Uranium	10
	1 ^{15, 19}
Volatile halocarbons	
Volatile halocarbons Chloroform	
Chloroform	0.1

Summary of WTS Performance Criteria (all concentrations in mg/L) Table 9.3:

- Notes:
 BOLD exceeds performance criteria

 1
 All concentrations in mg/L

 2
 LOR: limit of reporting

 3
 Sydney Water will introduce acceptance standards for a substance on a sub-system specific basis as

 determined by:
 - · how much the receiving system can transport and treat

how corroded the sub-system is how sewage treatment products will be used.

- 4. Discrete oil, fat or grease must not be discharged.
- 5. Where ammonia is present with other nitrogenous compounds, the amount of nitrogen in the ammonia is deducted from the Total nitrogen as measured by Total Kjeldahl Nitrogen, before calculating the charge for nitrogen.
- 6. Fluoride, phosphorus and nitrogen limits don't apply where the customer's sewerage system is connected to a sewage treatment plant that discharges to the ocean.
- Acceptance standards also apply to concentrations of ammonia, benzene, bromine, chlorine, cyanide, formaldehyde, petroleum hydrocarbons, sulphide and volatile halocarbons in discrete samples.
- 8. Sydney Water will determine acceptance standards for individual chlorinated phenolics on a catchment basis, following pollution reduction targets set by the DECCW NSW for the sewage treatment plant effluent. The concentration limit is a guide only and we may set lower limits for individual chlorinated phenolic compounds.
- 9. Sydney Water do not allow discharge from comfort air conditioning cooling towers and evaporative condensers using products containing hexavalent chromium (chromate) or organometallic algicides, if the blow down (or 'bleed-off') is connected to the sewer. Comfort cooling towers are defined as cooling towers dedicated to heating, ventilation, air-conditioning or refrigeration systems.
- 10. Cyanide is defined as labile cyanide amenable to alkaline chlorination. This includes free cyanide as well as those complex cyanides that are particularly dissociable, almost wholly, or in a large degree, and therefore potentially toxic in low concentrations.
- 11. Sydney Water will not consent to any discharge of organochlorine pesticides (including chlordane, dieldrin and heptachlor), or organophosphorus pesticides (including chlorpyrifos, diazinon and malathion) into the sewerage system.
- 12. The limit for lithium applies only to the Rouse Hill sewage catchment.
- 13. Where flammable and/or explosive substances may be present, the customer must demonstrate to us that there is no possibility of explosions or fires in the sewerage system. We will discuss limits and charges with individual customers, before a trade waste agreement is negotiated. The flammability of the discharge must never exceed five per cent of the Lower Explosive Limit (LEL) of hexane at 25 OC. In some cases a customer may be required to install an LEL meter.
- 14. Sydney Water will determine acceptance standards for total dissolved solids on a catchment-specific basis. A limit of 500 mg/L may apply to customers discharging to an inland sewage treatment plant or to a sewage treatment plant that is part of a designated reuse system. Acceptance standards will only apply to those customers discharging in excess of 100kg/d of total dissolved solids (TDS) or greater than one per cent of the total catchment TDS load (whichever is the lesser).
- 15. Analysis of volatile halocarbons must at a minimum include methylene chloride, chloroform, trichloroethylene and perchloroethylene.
- This substance is made up of several substances including benzene, toluene, ethylbenzene, (m+p)-xylene and o-xylene.
- 17. As at 1 July 2010, the limit for soluble BOD applies only to the Smithfield sewage and SPS 67 catchments, due to corrosion.
- 18. This is a guide only. Exact allowable levels are determined on a system-specific basis.
- 19. Charges will apply for total volatile halocarbons
- 20. Charges will apply for total petroleum hydrocarbons (flammable)



9.4 Details of Trialled WTS

The following WTS was trialled on site:

- Equalization/storage initial influent collection tanks to allow consistent loading to the WTP under variable conditions in the collection system and to optimize the size of the treatment system;
- Pre-treatment System comprising an oil/water separator to remove any free oily materials prior to treatment;
- Filtration System comprising;
 - Bag filters to remove particulate matter and to protect the downstream filtration processes;
 - Activated Carbon Filters to remove dissolved organics and some inorganic components;
 - Ancillary tanks, pumps, control and monitoring equipment.

A process flow diagram of the treatment system is presented in Figure 11.

Consideration should be given to the inclusion of influent tanks in the full scale system used for the remediation works given inflow rates from the fill layers are likely to vary across the site and over the duration of the works.

Influent into the system was restricted to 72 litres / min using a control valve. The flow rate adopted was equivalent to a contact time with the carbon filter of 8 minutes.

With the exception of arsenic, the effluent water generated by the system was generally compliant with ANZECC/ARMCANZ 2000 95% trigger values. The result of the trial indicated that arsenic concentrations increased following contact with the activated carbon filter. It was considered that this impact can readily be minimised by the selection of acid washed GAC filter for use in the operational WTP.

9.5 WTS Waste

A number of waste streams will be generated through use of a WTS and planning for the remediation works must consider the disposal requirements for these materials as follows.

- Free oil removed in the oil/water separator will require collection, most likely to be discharge to a 'drum' collection system. Any drums should be used and stored within a containment area to collect any spills. The collection system will be equipped with a level switch to advise the operator when the drums are full and require disposal, or systems shall be in place such that overflow of the drums does not occur;
- The filter bags in the sand bag filter system will have to be replaced and disposed of once the filters become filled with sediment. Testing of the filter bag was not conducted as part of the water treatment trial. It is possible that these filter bags are classified as a hazardous waste due to the presence of hydrocarbons and/or metals, and will require appropriate disposal to landfill; and
- The activated carbon filter media will become fully loaded over time and require disposal. The replacement frequency will depend on the contaminant load in the water for treatment. In adopting a sustainable approach to the works it is recommended that where possible the spent activated carbon media be returned to the supplier for regeneration, rather than disposal.



Appropriate disposal of wastes generated by the WTS may require sampling and analysis to determine the appropriate disposal location.



10 Considerations for Treatment of Contaminated Material

The revised remedial strategy for the site incorporates options for managing material containing exceedances of the acceptance criteria, including soil treatment by cement stabilisation (at Chullora) or bioremediation (on site).

One of two objectives exist for the inclusion of this option:

- Treating coal tar contaminated material such that a reduced waste classification may be achieved for disposal off site to landfill, in accordance with NSW EPA 'General Immobilisation of Contaminants in Waste – Coal tar Contaminated Waste From Former Gasworks Sites' approval number 2005/14 (IA 2005/14). Excavated material that requires off site disposal as Hazardous Waste will need to be treated in this manner. The Contractor may choose to similarly treat material that requires off site disposal as Restricted Solid Waste should their cost benefit analysis indicate worthwhile savings can be achieved in the project timeline or budget; or
- 2. Treating coal tar contaminated material such that it can be reused on site. This will require demonstration of the following for the material undergoing treatment by bioremediation on site compliance with the site specific total concentrations (as provided in Table 4.1 of the RAP) AND the material does not pose a risk to groundwater migrating off-site and is compliant with the site specific leachability criteria for the site.

Figures 12, 13 and **14** provide an estimate of the areas suited to treatment by cement stabilisation or bioremediation. Each figure relates to a particular soil depth interval. The areas and values provided in **Figures 12** to **14** should be viewed as indicative only and have been based on an assessment of the available historical data, including borelogs describing the prevalence of free tar impacts in soil and fill. Actual conditions encountered during bulk excavation of the site may vary.

Furthermore the volume of material suited to each treatment option after excavation will be heavily influenced by the methods adopted and strict implementation of site controls to prevent cross contamination.

10.1 Treatment Using Cement Stabilisation (Chullora)

In the benchscale immobilisation trial conducted by JBS (**Appendix A**) the results indicated that two of the three materials tested were capable of achieving the required UCS value of 1 MPa (NSW EPA 2005) with a minimum addition of 12.5% cement. The other parameters tested in these materials also showed full compliance with the requirements of IA 2005/14, and under this order would be suitable for off site disposal to landfill as General Solid Waste. The third material failed to meet the required UCS value even with 20% cement addition, and was assessed to be not stabilised. The failure was assumed to be related to the clay content of this material, which was noted to be 87% and well above the general range of 60% to 80% material considered suitable for cement stabilisation. As all other parameters in the failed material showed full compliance with the requirements of IA 2005/14, the stabilisation issue may be overcome by using an increased ratio of cement in the treatment process provided the mixing ratio provided in IA2004/14 Condition 1.3 is note exceeded.



It is noted that bioremediation will not be applicable to all excavated source materials from the site. Bioremediation should only be undertaken on material impacted by volatile contaminants, i.e. areas of the site impacted with heavy metal contamination cannot be bioremediated. Additionally where excavated material requires remediation of PAHs the decision to bioremediate should consider the form of the contamination and proportion of individual compounds present, noting that:

- Bioremediation of material impacted by free tar is unlikely to be practicable; and
- Bioremediation is applicable to material impacted with lighter end PAHs such as naphthalene, while bioremediation of the heavy end PAHs may not at all be possible in a reasonable timeframe or may require the application of additives for breakdown.

The effectiveness of the bioremediation process will also be dependent on the layout of the bioremediation area and the frequency of turning adopted.

10.3 Off Site Treatment

Given the proximity of the Macdonaldtown site to residents, and likely space restrictions during excavation of contaminated soils, an alternate site has been identified by RailCorp for ex-situ treatment of soils. Approval is being sought for treatment to occur on an approximate 2 ha parcel of land contained within the RailCorp owned Chullora Railway Workshops, located on Worth Street, Chullora, NSW. A plan showing the likely maximum area to be made available for treatment works is provided as **Figure 15.** Material treated off site must be disposed to an appropriately licensed facility.

10.4 Storage Volume

Storage volumes for Macdonaldtown have not been estimated given the inherent variability in how remedial excavations may be staged.

The volume of material capable of being stored and or treated within the designated area at Chullora will depend on several project specific factors including the treatment method to be adopted, the area required for truck loading areas, plant and supplies, and the size of the allocated area for treatment.

To aid with planning for the site **Table 10.1** provides indicative estimates of volumes of material that can be handled on the Chullora site. Each process has been assessed individually *i.e.* volume of soil if all treatment area use for either storage, curing of stabilised material or bioremediation windrows. The values assume minimal area only is required for truck loading, supply and plant storage. The approximate volume of material treatable is likely to be easily estimated by determining what percentage of the site is to be used for each task.

Process	Area Occupied	Indicative Volume capable of storage on site at one time	Comments/ Assumptions					
Storage	0.5 ha ¹	8,250 m ³	Stockpile max 5 m high, 10m wide, in 5 rows running lengthwise across area					
Treated Stabilised soil – layed out for curing period	0.5 ha ¹	2,500 m ³	Treated soil placed in 0.5m high blocks for curing					
Soil in wind rows for bioremediation	0.5 ha ¹	1250 m ³ (storage on site at any one time)	Soil for bioremediation place in 1 m high windrows in 8 rows running lengthwise across site.					
		approximately 50 m ³ uncovered (i.e. being treated at any one	Assumes that the maximum number of rows that can be uncovered at any one time should be consistent with the requirements of the air quality					

Table 10..1 – Estimated Volume of Material Capable of being stored in Chullora Treatment Area



Notes: 1 actual volumes will depend of dimensions of available area

10.5 **Treatment Rates**

Table 10.2 presents possible treatment rates achievable based on discussions with remediation contractors. It is noted that the rate at which material can be treated either on the Macdonaldtown or Chullora sites, will be entirely dependent the manner in which treatment is undertaken, including the method adopted, plant utilised and site configuration.

Process	Likely treatment rate	Comments/ Assumptions
Treatment by Cement Stabilisation	1000 T per day ¹ , can be streamlined to achieve up to 1400 T per day	Daily excavation rates are unlikely to match treatment of 1000 tonnes per day. Considerations should be given to either use of smaller mixing plants in line with excavation rates, or if large daily treatment volumes are planned whether sufficient space exists for pre-treatment stockpiling and post treatment curing
Treatment by bioremediation	500 m ³ per 3 weeks (average) ^{2.3}	Bioremediation may not be applicable to all contaminants requiring remediation

Notes: 1 achieved using semi-trailer sized mixing plant

2 assumes maximum volume possible stored on treated site, staggered such that all 500 m³ is subject to daily exposure, with no more than 150m² uncovered at any one time. 3 treatment rates will vary if bioremediation to be completed within tented enclosure.



11 Considerations for Material Disposal

Table 8.1 includes summary of the range of materials likely to require management duringthe remediation program. Materials anticipated to be classed as Restricted Solid Waste orGeneral Solid Waste will be suitable for direct disposal to appropriately licensed landfills.

As no landfills within NSW are currently licensed to dispose of Hazardous Waste, should these materials require off site disposal, the following will be required:

- treatment to achieve a lower Waste Classification prior to disposal (to a landfill licensed to receive treated material of that class of treated material); or
- delivery to an off-site facility licensed to store the material, prior to treatment for disposal to an appropriately licensed landfill.

The requirements of treatment and disposal to landfill are discussed in the Sections below.

11.1 Treatment Using Bioremediation

The objective of pre-treatment of materials classed as 'Hazardous Waste' by bioremediation would be to reduce the classification to Restricted Solid Waste or General Solid Waste. As the method should reduce contaminant mass, successful treatment should allow for unrestricted disposal of the material at an appropriately licensed landfill.

11.2 Treatment Using Cement Stabilisation

If material is to be pre-treated using cement stabilisation, the works will require completion in accordance with the DECC '*Immobilisation Technical Note 1 - Process Equipment for Treatment of Contaminated Soil and Sludge Waste.*' A copy of this document is provided in **Appendix E**.

If the material is to be disposed to landfill following cement stabilisation, disposal must be in accordance with the requirements of NSW EPA IA 2005/14, specifically that:

- The treated waste can only be disposed of at a landfill that has 'currently operating leachate management systems' and which are licensed by DECCW to receive that particular class of waste;
- The landfill licence allows the site to receive 'waste subject to immobilisation approvals with this type of disposal restriction';
- If the total concentration in the waste exceeds either the Specific Contaminant Concentration (SCC)2 or SCC3 values (Waste Classification Guidelines 2009), the landfill monitors leachate and groundwater for PAHs or B(a)P; and
- The landfill must be advised in writing that the material has been treated and classified in accordance with the requirements of IA #2005/14.



If material is to be treated by cement stabilisation for disposal to landfill in accordance with IA2005/14, all the requirements of this general immobilisation approval must be satisfied, including, but not limited to:

- Ensuring the proposed location of cement stabilisation is lawfully able to treat the waste, and in consultation with RailCorp, obtaining an permits or licences deemed necessary for the works;
- Providing notification to the DECCW of the intention to treat and dispose of material in accordance with IA2005/14 at least 28 days prior to the commencement of these works;
- Use of calcium or magnesium oxide based cement;
- Ratio of reagent to untreated waste must not exceed 2:1; and
- The UCS of the treated waste must be measured in accordance with NSW Roads and Traffic Authority (RTA) Test Method T131 of T116. The use of equivalent methods will require receipt of written approval from DECCW.

Furthermore if material is to be treated by cement stabilisation, the works will require completion in accordance with the DECC Immobilisation Technical Note 1 - *Process Equipment for Treatment of Contaminated Soil and Sludge Waste.* A copy of this document is provided as **Appendix E.**

It is noted that the conditions listed in IA 2005/14 preclude use of the approval on materials containing Total PAH concentrations in excess of 13 000 mg/kg or B(a)P concentrations in excess of 500 mg/kg. Review of historical site data indicates that the total concentrations in three samples analysed from the site exceeded these limits, specifically:

- free tar impacted clay at MG08/1.5 m (Total PAH 15 237 mg/kg);
- two pipe contents samples labelled as 'RP' and 'Service Pipe' and described as coal tar material sampled from a pipe (Total PAH of 20 890 mg/kg and 26 805.3 mg/kg);

The last two samples are likely to represent the concentrations in free tar present in residual underground infrastructure. This material is likely to be disposed off site as liquid waste (as described in **Section 8**), and is unlikely to require treatment for stabilisation.

The sample collected from MG08 represents clay material heavily impacted by free tar. Review of the borelog for this location indicates that the material sampled at 1.5 m was a thin band of impacted soil. When considered in isolation this thin band of soil is unsuitable for treatment by cement stabilisation, and may require transfer to a licensed off-site storage site (discussed further in **Section 11**). However, given that the great majority of all other soil samples analysed from the site contained total PAH concentrations less than half of the MG08/1.5 m concentration, it is considered unlikely that when this area is bulk excavated and handled during treatment that PAH concentrations consistently in excess of 13,000 mg/kg would be detected.

10.2 Treatment Using Bioremediation

No bioremediation trials were undertaken as part of the pre-remedial investigations. However bioremediation has been demonstrated as an effective remediation technology on former gasworks sites when used in conjunction with other technologies. Use of bioremediation wherever possible, as an auxiliary process in the proposed works, has the potential to realise significant cost savings and reductions in the volume of material disposed to landfill.



12 Indicative Program

The anticipated program for the remediation works is summarised in **Table 12.1.** The proposed program is based on the following assumptions:

- Excavation of 350 m³ of soil on average per day from the site to the designated Chullora Treatment Area or directly to landfill over an initial three month period;
- Treatment of excavated soil as received, and for a further month after all remedial excavations completed at Macdonaldtown; and
- Site reinstatement occurring once validation of excavations completed, and continuing for a further three months after completion of treatment works.

The actual time required for completion will be dependent on the specifics of the remedial option adopted and time required for any additional administrative requirements specified by RailCorp.





Table 12.1 – Anticipated Program of Remediation Works

	Month Number																									
Stage	1	7	£	4	n	n	9	2	ø	6	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25
Pre-Remedial works ¹																										
Planning and site Establishment																										
Excavation of Contaminated soil (up to 23 000 m ³)																										
Validation of excavations, review of interim results by RailCorp + Site Auditor																										
Disposal to landfill of untreated material (estimate up to 5300 m3)																										
Treatment of soils by stabilization (assume up to 9800 m ³)																										
Treatment of soils by bioremediation (assume 50% of up to 5300 m ³)																										
Reinstatement to pre remediation levels ³																										
Post remediation studies and monitoring ³																										

Notes: 1 includes time for site preparation of project site specific management plans including occupation health and safety 2 includes all heritage surveys, vegetations management plans and structural surveys required 3 includes landscaping and revegetation, completion of validation reports and post remediation EMP for the site

4 duration of groundwater monitoring to be determined in the post remediation EMP for the site



13 Limitations

This report has been prepared for use by the client who commissioned the works in accordance with the project brief only and has been based in part on information obtained from other parties. The advice herein relates only to this project and all results conclusions and recommendations made should be reviewed by a competent person with experience in environmental investigations, before being used for any other purpose.

JBS Environmental Pty Ltd accepts no liability for use or interpretation by any person or body other than the client. This report should not be reproduced without prior approval by the client, or amended in any way without prior approval by JBS Environmental Pty Ltd, and should not be relied upon by other parties, who should make their own enquires.

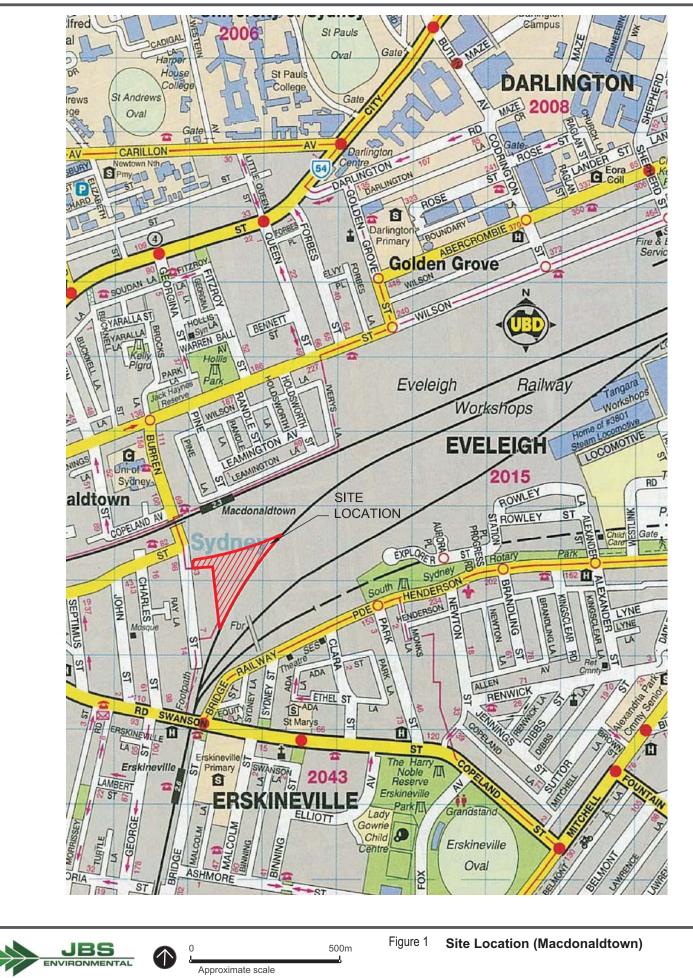
Sampling and chemical analysis of environmental media is based on appropriate guidance documents made and approved by the relevant regulatory authorities. Conclusions arising from the review and assessment of environmental data are based on the sampling and analysis considered appropriate based on the regulatory requirements and site history, not on sampling and analysis of all media at all locations for all potential contaminants.

Changes to the subsurface conditions may occur subsequent to the investigations described herein, through natural processes or through the intentional or accidental addition of contaminants. The conclusions and recommendations reached in this report are based on the information obtained at the time of the investigations.

This report does not provide a complete assessment of the environmental status of the site, and it is limited to the scope defined herein. Should information become available regarding conditions at the site including previously unknown sources of contamination, JBS Environmental Pty Ltd reserves the right to review the report in the context of the additional information.



Figures



CH2M Hill (2007) Note- All locations shown are approximate only

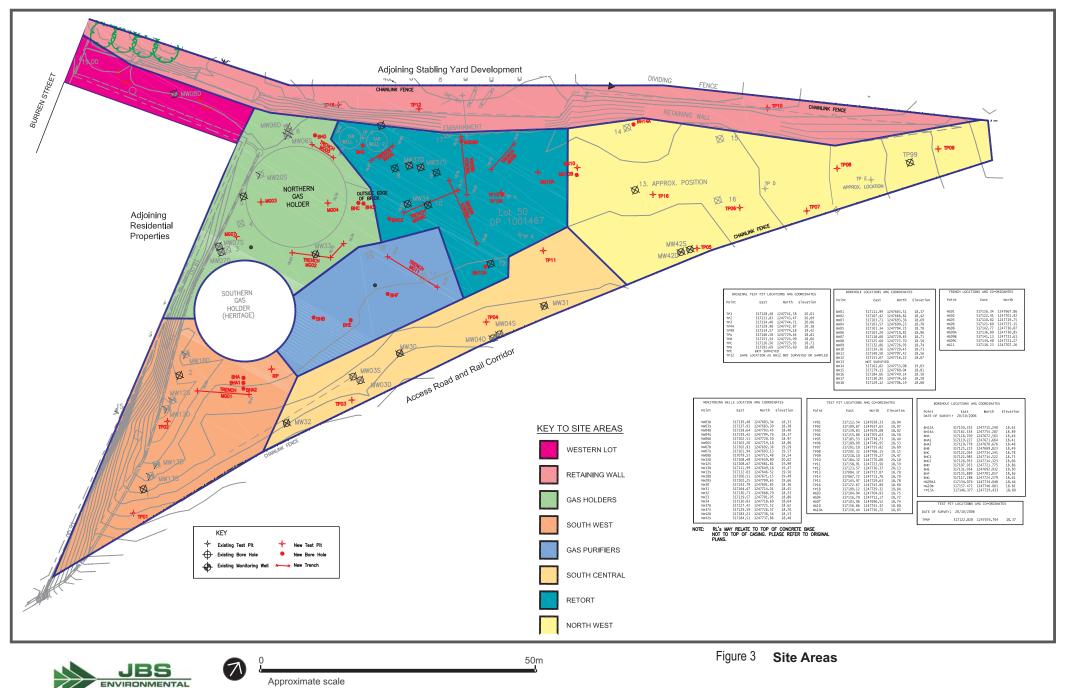




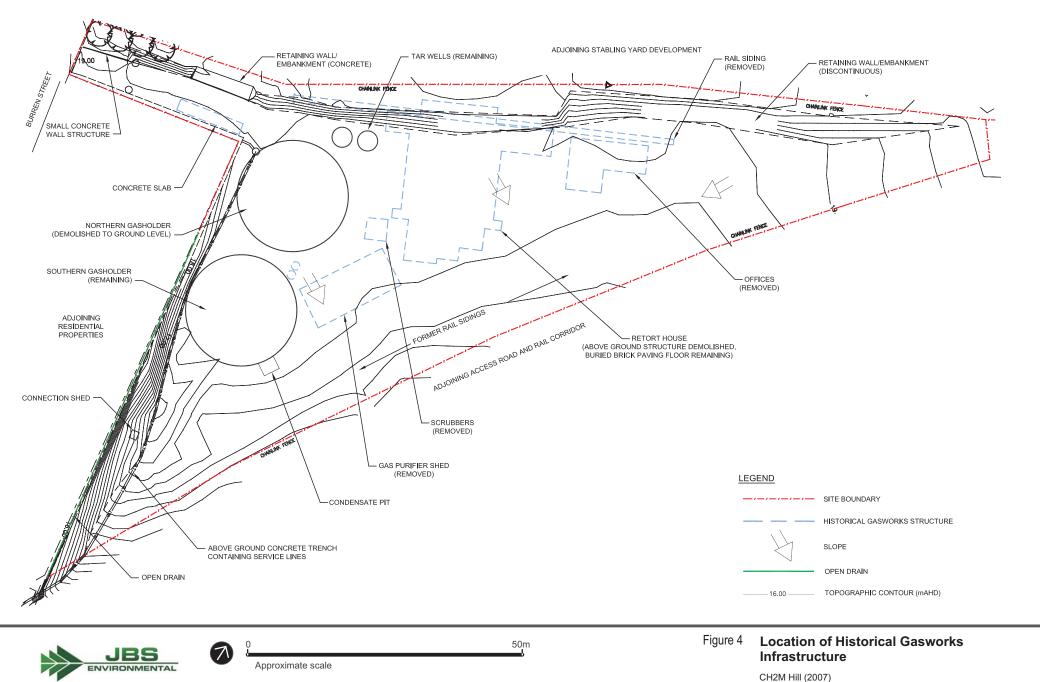
Approximate scale

Figure 2 Current Macdonaldtown Site Plan

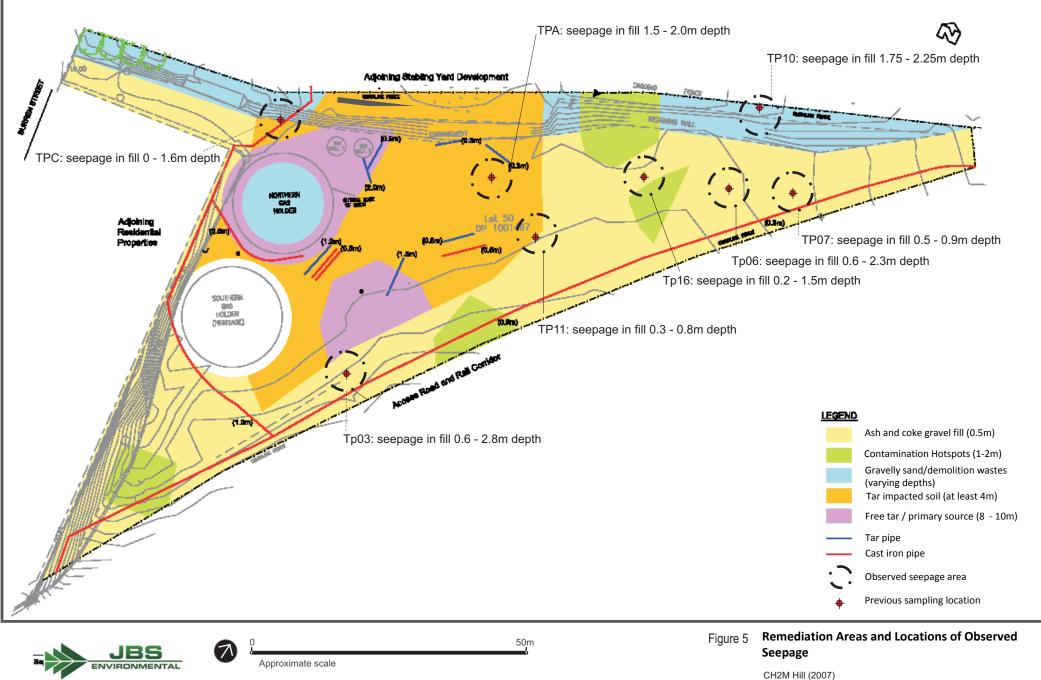
CH2M Hill (2007) Note- All locations shown are approximate only



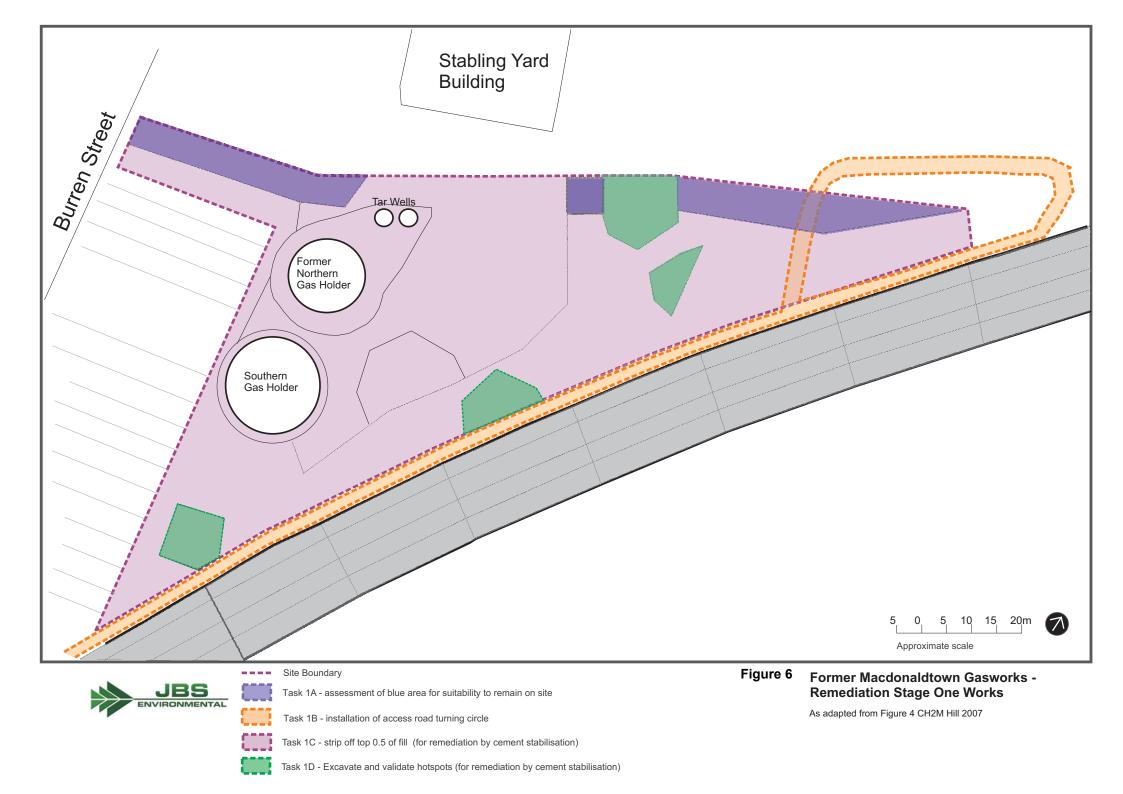
CH2M Hill (2007) Note- All locations shown are approximate only

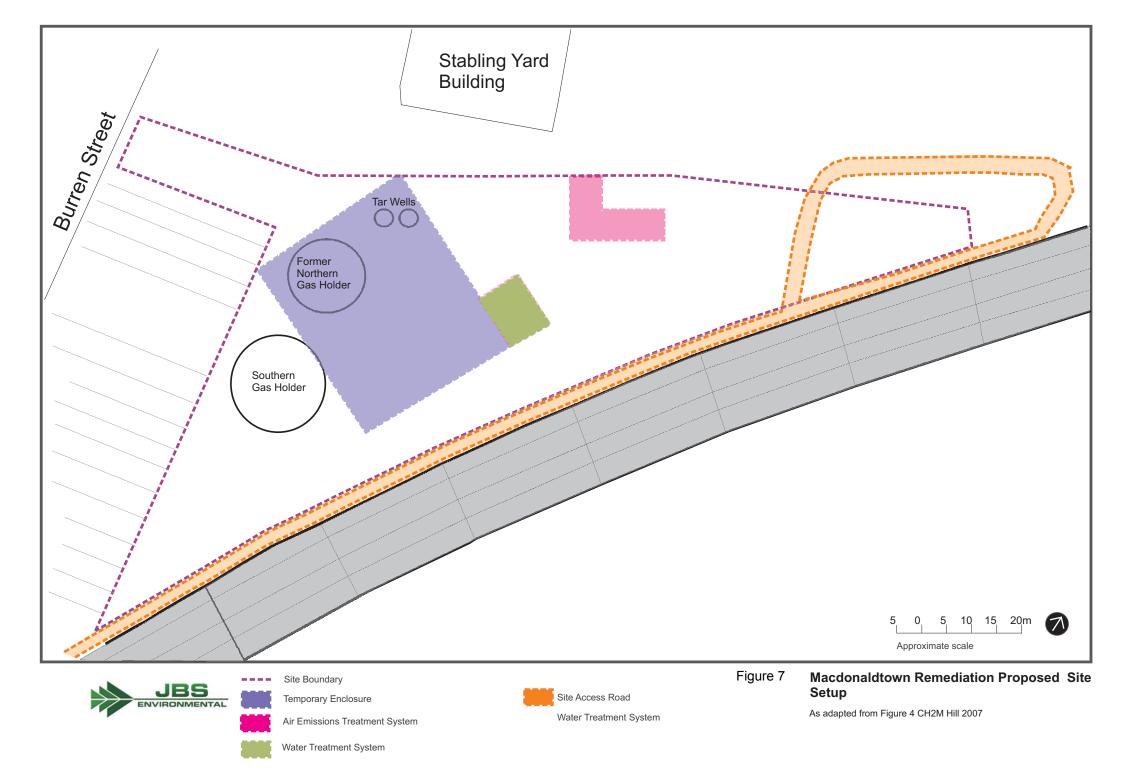


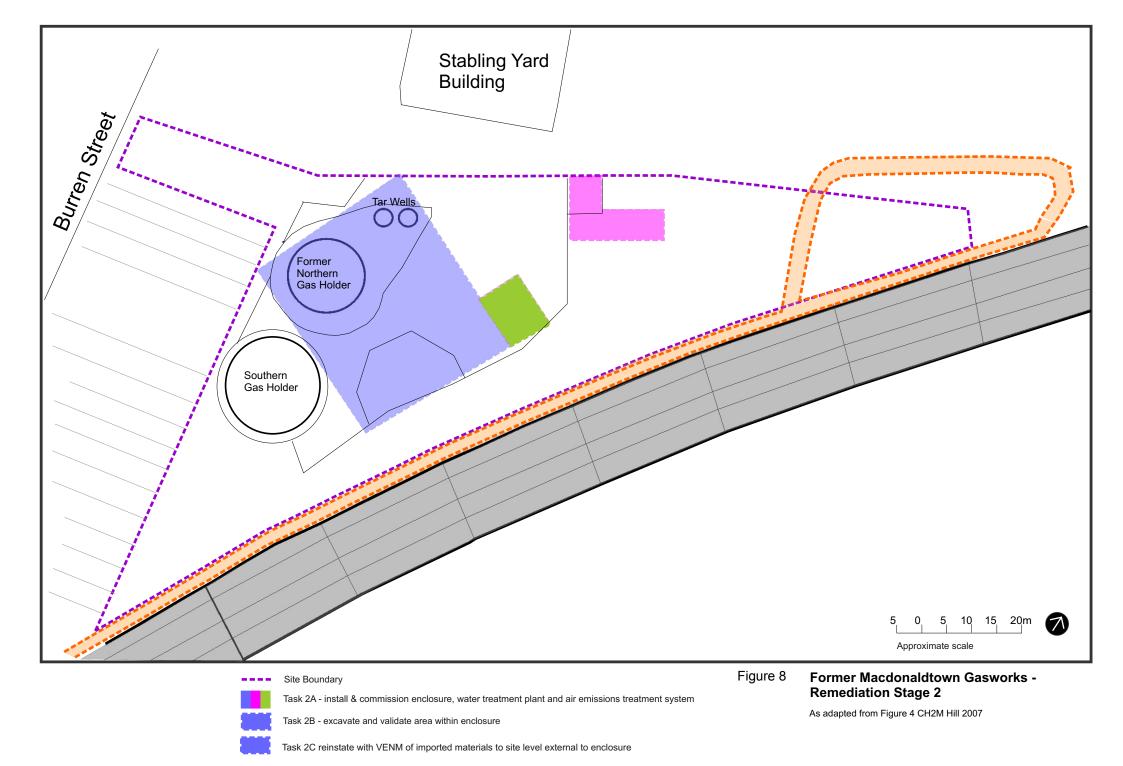
Note- All locations shown are approximate only

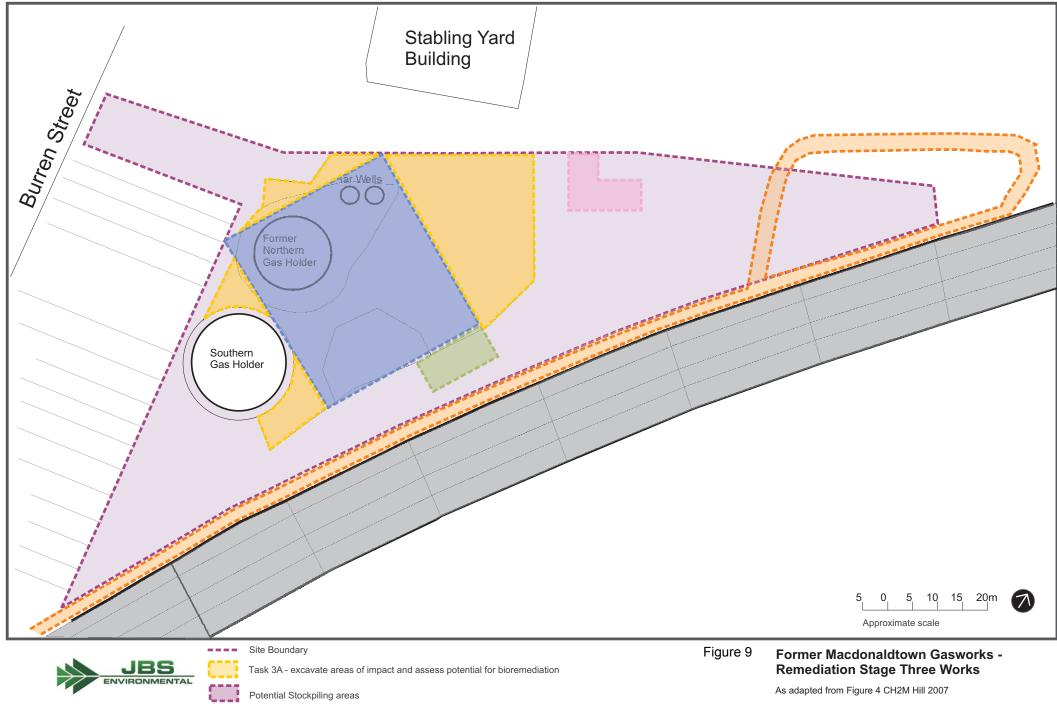


Note- All locations shown are approximate only



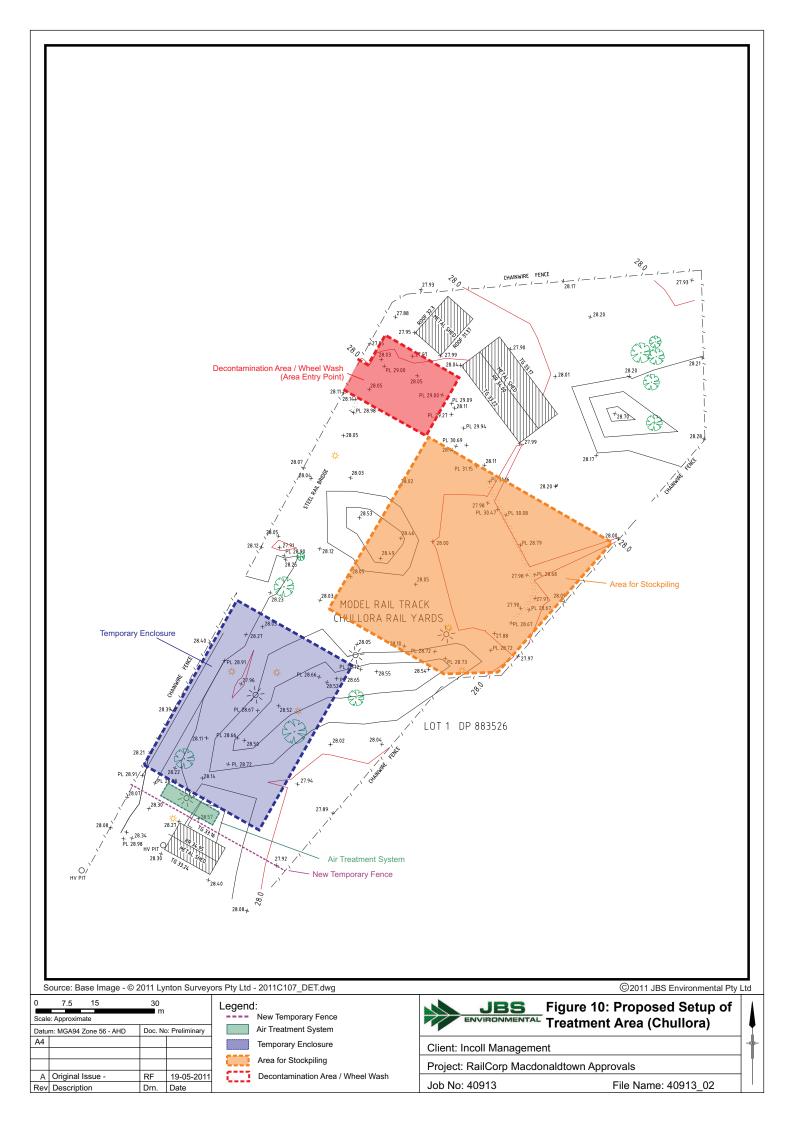




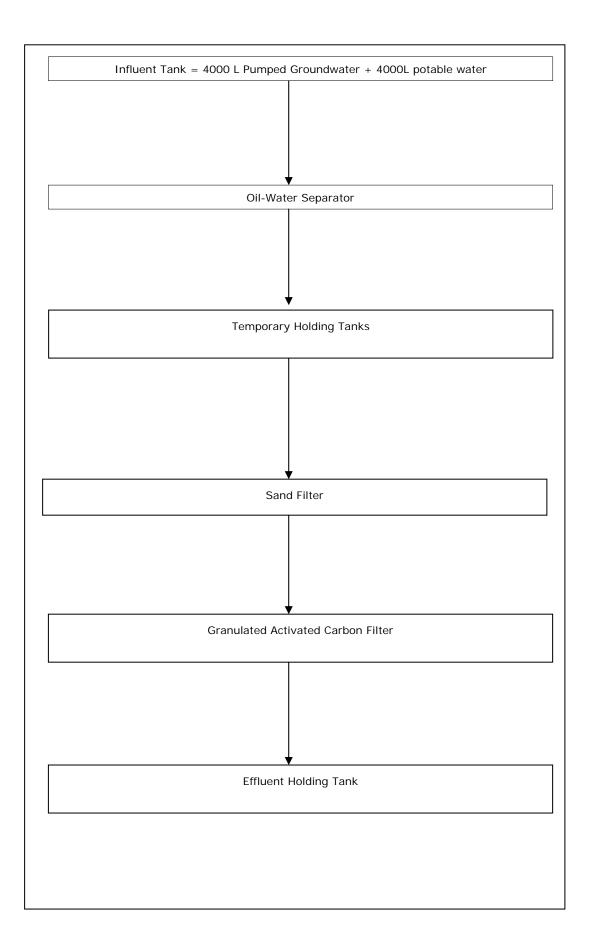


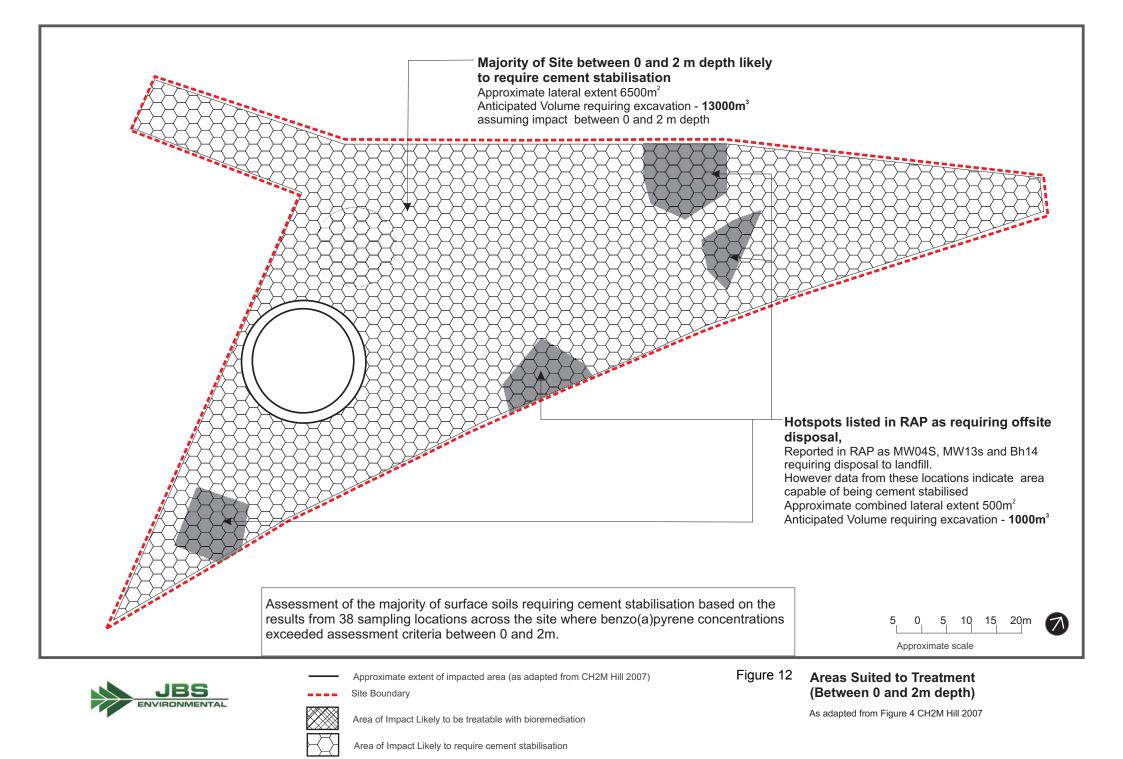
Potential Stockpiling areas

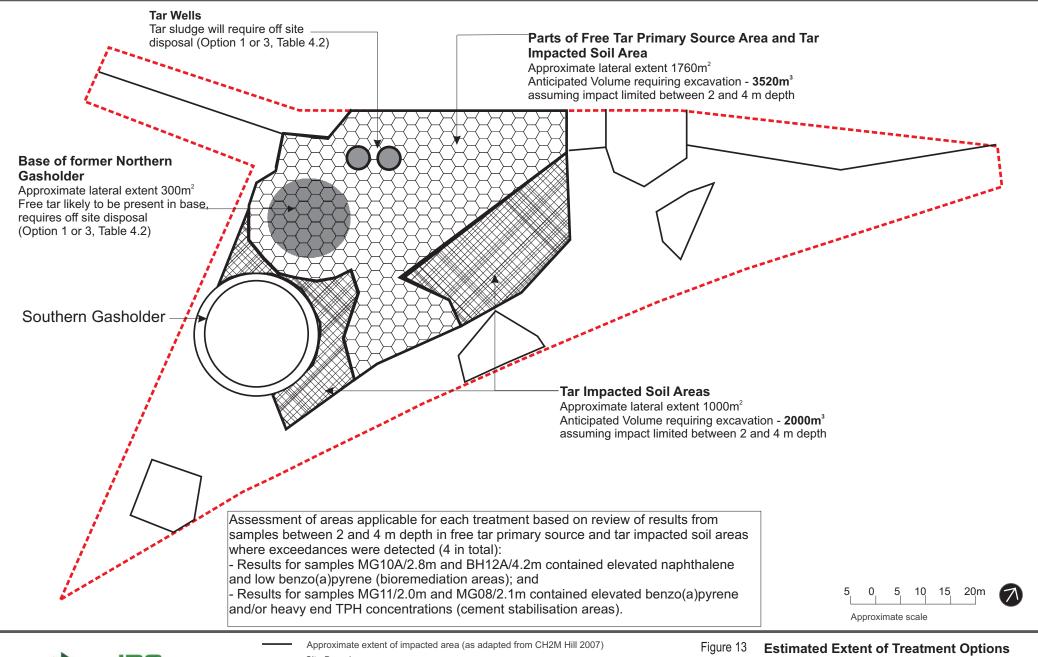
As adapted from Figure 4 CH2M Hill 2007











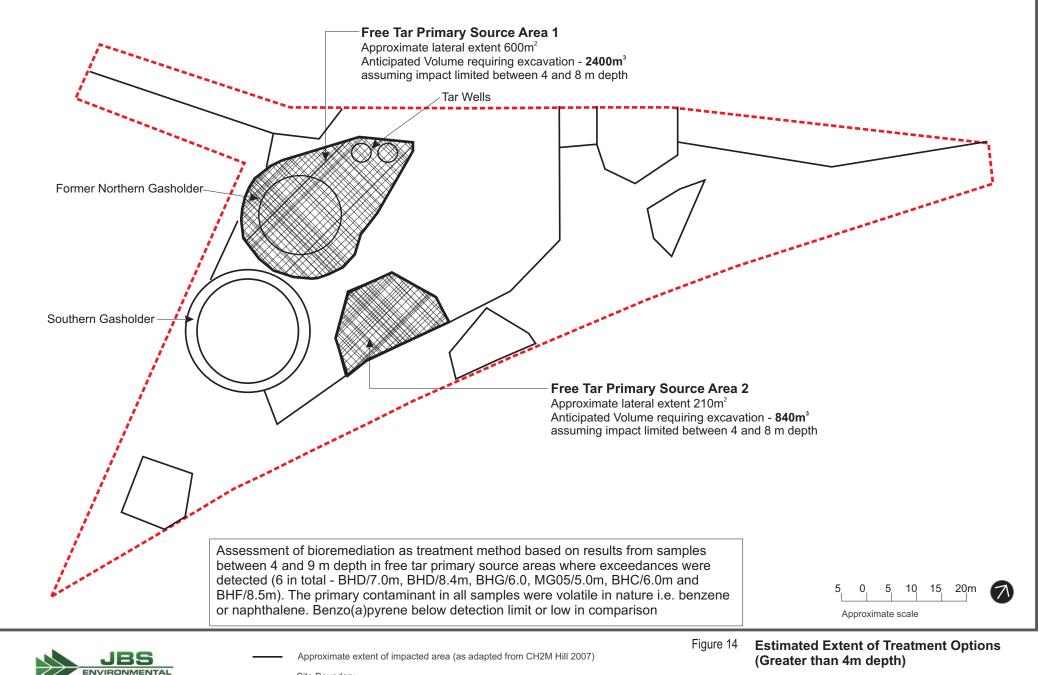


Site Boundary Area of Impact Likely to be Subject to Treatment or Management by Options 2 or 3 in Table 4.2 (bioremediation)

Area of Impact Likely to be Subject to Treatment or Management by Option 3 in Table 4.2 (cement stabilisation)

(Between 2 and 4m depth)

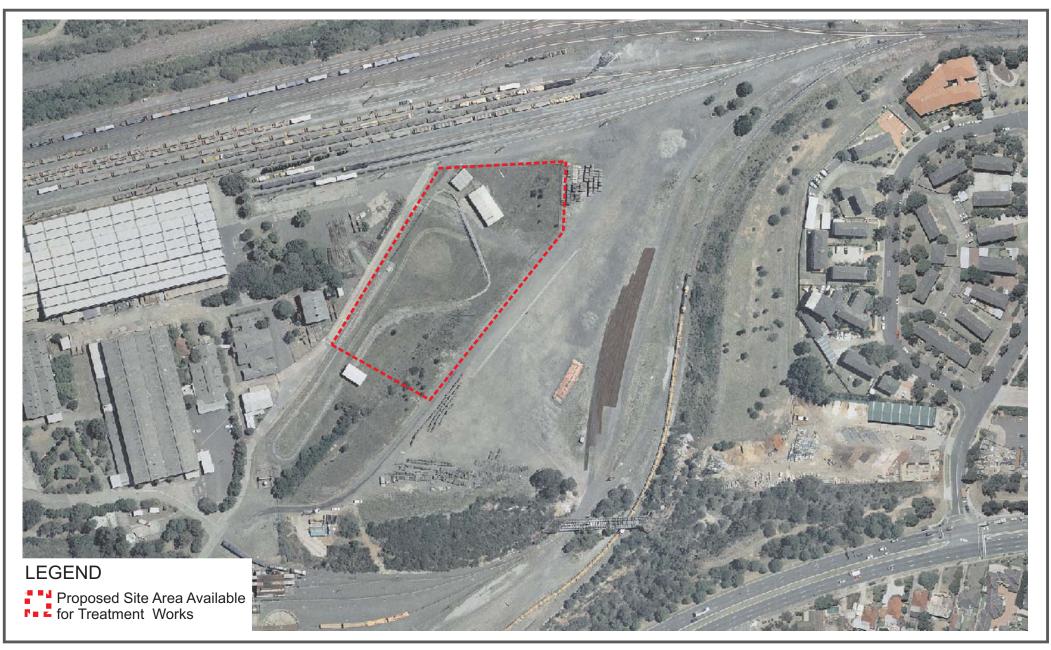




Site Boundary

Area of Impact Likely to be subject to Treatment or Remediation in accordance with Option 2 or 3 in Table 4.2

As adapted from Figure 4 CH2M Hill 2007





50m

Figure 15 Chullora Treatment Area Site Plan

Department of Lands (2010) Note- All locations shown are approximate only



Tables

Table A: Revised Remedial Strategy – Areas, Extents and Likely Treatment Requirements

		E			Primary Treatment		
Site Area	Material Type	Extent of Impact	Expected Quantity (m ³)	Contaminants (Maximum reported concentration)	Treatment Options (1,2, 3 or 4 from Table 4.2)	Anticipated Waste Classification Subsequent to Treatment / Immobilisation Approval	Licencing Req
	Tar Sludge	Contained within Base of Gasholder	320	Benzo(a)pyrene (595 mg/kg) Total PAHs (26 805.3 mg/kg) TPH C6-C9 (3770 mg/kg) TPH C10-C14 (1 180 000 mg/kg) Benzene (814 mg/kg) Ethylbenzene (254 mg/kg) Total Xylenes (3170 mg/kg)	Option 1: Liquid Waste Disposal	-	No premises ba Licenced Liquid treatment/dispc
	Impacted Water	Contained within Gasholder	640	Naphthalene (1 460 µg/L) Total PAHs (1 6774 µg/L) TPH C6-C9 (2170 µg/L) TPH C10-C14 (9495 µg/L) Benzene (704 µg/L) Ethylbenzene (213 µg/L) Toluene (117 µg/L) Total Xylenes (417 µg/L))	Option 1: Liquid Waste Disposal; or disposal to sewer or on site reuse	Suitable for disposal to sewer or on site beneficial reuse	Disposal to sew
nasnoider	Soil / fill impacted by free tar	Base annulus and proximate soils (within pink shaded area on Figure 5 to a depth of 8m-10m)	2100	Benzo(a)pyrene (17.6 mg/kg) Total PAHs (1906.4 mg/kg) TPH C6-C9 (559 mg/kg) TPH C10-C14 (8760 mg/kg) Benzene (5.4 mg/kg) Ethylbenzene (40.8 mg/kg) Total Xylenes (246.7 mg/kg)	Options 2 or 3: Treatment by bioremediation treatment for onsite reuse or landfill disposal OR Treatment by stabilisation to apply NSW DEC immobilisation approval (Approval #2005/14)	Restricted Solid Waste or General Solid Waste (may also be assessed for suitability for on site reuse)	For treatment - accordance with #2005/14), Dev immobilisation a For off-site disp - be licensed th - have an opera - monitor leacha
	Demolition			Benzo(a)pyrene (17.6 mg/kg) Total PAHs (1906.4 mg/kg) TPH C6-C9 (559 mg/kg) TPH C10-C14 (8760 mg/kg) Benzene (5.4 mg/kg) Ethylbenzene (40.8) Total Xylenes (246.7) Asbestos present		Hazardous Waste (free tar)	No premises ba Licensed Liquid treatment/dispo
			1900			Special Waste (asbestos)	For site remova works must be For off-site disp waste
					General Solid Waste (Non-Putrescible)	For off-site disp Solid Waste (no For recycling – recovery catego	
	Impacted Soil	Lateral extent of orange shaded area on Figure 5 to a depth of at least 4m	9225	Benzo(a)pyrene (444 mg/kg) Total PAHs (15 237.6 mg/kg) TPH C6-C9 (51 mg/kg) TPH C10-C14 (435 100 mg/kg) Benzene (0.3 mg/kg) Ethylbenzene (12.3 mg/kg) Total Xylenes (10.6 mg/kg)		Restricted Solid Waste or General Solid Waste (may also be assessed for suitability for on site reuse)	For treatment -
Retort and Surrounding Former Gasworks Source Areas	Impacted Natural Soil	Lateral extent of pink shaded area on Figure 5 in the vicinity of boreholes BHE and BHF to a depth of 8m-10m	2375	Benzo(a) pyrene (13.9 mg/kg) Total PAHs (515.6 mg/kg) TPH C6-C9 (228 mg/kg) TPH C10-C14 (5350 mg/kg) Benzene (20 mg/kg) Ethylbenzene (8.3 mg/kg) Total Xylenes (94.9 mg/kg)	Options 2 or 3: Treatment by bioremediation treatment for onsite reuse or landfill disposal OR Treatment by stabilisation to apply NSW DEC immobilisation approval (Approval #2005/14)	Restricted Solid Waste or General Solid Waste (may also be assessed for suitability for on site reuse)	- be licensed to assessed as;
hotspot at	Contamination	Lateral extent of green shaded area on Figure 5 to a depth of 1m-2m	115	Benzo(a)pyrene (39.4 mg/kg) Total PAHs (425.1 mg/kg) TPH C6-C9 (166 mg/kg) TPH C10-C14 (7640 mg/kg) Benzene (3.1 mg/kg) Ethylbenzene (6.4 mg/kg) Total Xylenes (61.2 mg/kg)		Restricted Solid Waste or General Solid Waste (may also be assessed for suitability for on site reuse)	- have an opera - monitor leacha
Existing Site Surfaces		Lateral extent of shaded area on Figure 5 to a depth of at least 0.5 m	2950	Benzo(a)pyrene (39 mg/kg at BH13) Total PAHs (413.2mg/kg at BH13) TPH C10-C14 (7100 mg/kg at BH13) Lead (2140 mg/kg at VP01 surface)	Options 2 or 3: Treatment by bioremediation treatment for onsite reuse or landfill disposal OR Treatment by stabilisation to apply NSW DEC immobilisation approval (Approval #2005/14)	Restricted Solid Waste or General Solid Waste (may also be assessed for suitability for on site reuse)	For treatment - accordance with #2005/14), Dev immobilisation a For off-site disp - be licensed to assessed as; - have an opera - monitor leacha
Retaining Wall	General Fill and demolition waste	Entire Northern boundary (shaded blue on Figure 5)		Benzo(a)pyrene 150 mg/kg) Total PAHs (2472.4 mg/kg) Benzene (15.0 mg/kg) Total Xylenes (1.7 mg/kg)	Landfill disposal, Beneficial Reuse or Recycling	-	For removal wo must be underta For off-site disp Solid Waste (no



quirements

based licence required for on-site handling id Waste contractor required for transport and ultimate sposal of free tar

ewer requires trade waste agreement with Sydney Water - Cement stabilisation may be undertaken in ith NSW DEC general immobilisation approval (Approval Deviations from this method may require a specific n approval to be obtained from DECCW sposal – landfill receiving treated material must: the class of waste the treated material is assessed as; erational leachate collection system chate and groundwater on site for B(a)P and Total PAHs based licence required for on-site handling id Waste contractor required for transport and ultimate posal of free tar val works – if more than 10m² of ACM encountered, be undertaken by a licencedAS1 contractor sposal – landfill must be licensed to received asbestos sposal – landfill must be licensed to received General non-putrescible) – premises must be licenced for appropriate waste gory

t - Cement stabilisation may be undertaken in vith NSW DEC general immobilisation approval (Approval Deviations from this method may require a specific n approval to be obtained from DECCW isposal – landfill receiving treated material must: I to receive the class of waste the treated material is

erational leachate collection system chate and groundwater on site for B(a)P and Total PAHs

t - Cement stabilisation may be undertaken in with NSW DEC general immobilisation approval (Approval Deviations from this method may require a specific n approval to be obtained from DECCW sposal – landfill receiving treated material must: to accept the class of waste the treated material is

erational leachate collection system chate and groundwater on site for B(a)P and Total PAHs works – if more than 10 m² of ACM encountered, works ertaken by a licenced AS1 contractor sposal – landfill must be licensed to received General non-putrecible) and /or Special Waste (Asbestos Waste)

					Primary Treatment		
Site Area	Material Type	Extent of Impact	Expected Quantity (m ³)	Contaminants (Maximum reported concentration)	Treatment Options (1,2, 3 or 4 from Table 4.2)	Anticipated Waste Classification Subsequent to Treatment / Immobilisation Approval	Licencing Requ
							as appropriate For recycling – p recovery catego
Hotspots	at locations	Lateral extent shown as green shaded area on Figure 5 to a depth of 1 m-2 m	340	At BH14: Benzo(a)pyrene (5 mg/kg) Benzene (4.6 mg/kg) Total Xylenes (48 mg/kg) At MW13: Naphthalene Benzo(a)pyrene (34.9 mg/kg) Total PAHs (346 mg/kg) TPH C10-C14 (6444 mg/kg) At MW04: Benzene (4 mg/kg)	Options 2 or 3: Treatment by bioremediation treatment for onsite reuse or landfill disposal OR Treatment by stabilisation or Stabilisation to apply NSW DEC immobilisation approval (Approval #2005/14)	Restricted Solid Waste or General Solid Waste	For treatment - accordance with #2005/14), Dev immobilisation a For off-site dispo - be licensed to material is asses - have an operat - monitor leacha
	Old Gasworks Pipes	varied	Unknown	Benzo(a)pyrene (595 mg/kg) Total PAHs (26 805.3 mg/kg) TPH C6-C9 (3770 mg/kg) TPH C10-C14 (1180 000 mg/kg) Benzene (814 mg/kg) Ethylbenzene (254 mg/kg) Total Xylenes (3170 mg/kg)	Option 1: Separate landfill disposal of empty pipework and tar contents	-	No premises bas Licenced Liquid \ treatment/dispo
Site Wide	Fill and natural soil materials	NA	Unknown	unknown	Option 2 or 3: Treatment by bioremediation treatment for onsite reuse or landfill disposal OR Treatment by stabilisation or Stabilisation to apply NSW DEC immobilisation approval (Approval #2005/14)	Restricted Solid Waste or General Solid Waste	For treatment - accordance with #2005/14), Dev immobilisation a For off-site dispo - be licensed to assessed as; - have an operat - monitor leacha
	Residual tar sources – subsequent to source removal	Unknown	Unknown	Unknown	Option 2 or 3: Treatment by bioremediation treatment for onsite reuse or landfill disposal OR Treatment by stabilisation or Stabilisation to apply NSW DEC immobilisation approval (Approval #2005/14)	Restricted Solid Waste or General Solid Waste	For treatment - accordance with #2005/14), Dev immobilisation a For off-site dispo - be licensed to assessed as; - have an operat - monitor leacha



quirements

e – premises must be licenced for appropriate waste gory

 Cement stabilisation may be undertaken in ith NSW DEC general immobilisation approval (Approval Deviations from this method may require a specific in approval to be obtained from DECCW sposal – landfill receiving treated material must: to receive to accept the class of waste the treated sessed as;
 erational leachate collection system

chate and groundwater on site for B(a)P and Total PAHs

based licence required for on-site handling id Waste contractor required for transport and ultimate posal of free tar

It - Cement stabilisation may be undertaken in ith NSW DEC general immobilisation approval (Approval Deviations from this method may require a specific in approval to be obtained from DECCW sposal – landfill receiving treated material must: to accept the class of waste the treated material is

erational leachate collection system <u>chate and groundwater on site for B(a)P and Total PAHs</u> t - Cement stabilisation may be undertaken in with NSW DEC general immobilisation approval (Approval Deviations from this method may require a specific n approval to be obtained from DECCW sposal – landfill receiving treated material must: to accept the class of waste the treated material is

erational leachate collection system chate and groundwater on site for B(a)P and Total PAHs



Appendix A Pre-Remedial Investigations



A1 Pre-Remedial Investigations -Methods

The short list of most remedial options applicable to the site has been determined as follows:

- Excavation and off-site disposal to landfill of all material containing contaminant concentrations above the site acceptance criteria;
- Excavation and treatment of all possible material containing contaminants concentrations above the site acceptance criteria, for either:
 - off-site disposal to landfill; or
 - on site reuse.
- Assessment of leachability and contaminant source zones on the site. Excavation
 of all accessible source zone materials and treatment, as required, for off-site site
 disposal to landfill and/or on-site reuse. Capping and on-site management of
 impacted areas assessed to be either:
 - inaccessible due to heritage or geotechnical constraints; and/or
 - outside of source zones and not posing an unacceptable risk to groundwater migrating offsite.

Based on the available dataset there were several items that required additional information to enable design of a remedial strategy. The following gaps /areas requiring resolution were identified:

- Can tar impacted soils be stabilised, and if so what pre-treatment works may be required? Specifically, it is noted that while the RAP recommends that much of the impacted material be remediated by stabilisation or thermal treatment for off-site treatment, no assessment has been undertaken on the potential applicability of either treatment method. As the stabilisation of soil is likely to be the more cost effective treatment method of the two specified, it was considered prudent to run a bench scale trial of the required methodology, in accordance with NSW EPA (2005), to verify its suitability for application at the site;
- If it can be demonstrated that tar impacted soils can be stabilised, what is the suitability the stabilised material to be retained on site? Specifically what are the maximum leachable concentrations of the contaminants of concern (COC) in soil that will not have an adverse impact on the surrounding environment? Results of the bench scale trial could be used to assess the maximum leachable concentrations in soils (treated or untreated) that can be retained on site. These concentrations, incorporating factors of dilution that may occur as leached contamination mixes with groundwater, and fate and transport modelling on the mixed water, could be incorporated into the remedial strategy as clean-up criteria;
- What are the quantities of water likely to be generated during dewatering of the site for remediation? Previous investigations of the site have reported that contaminated soil is present at depths below the level of the perched water table in fill. Additionally, significant volumes of seepage have occurred in test pits excavated into the site. Efficient completion of the proposed remediation will therefore, require dewatering of fill materials in areas requiring remediation;
- What on site technologies will be suited to treatment of water generated during dewatering? Water samples collected from the perched water table in fill were reported to contain elevated concentrations of TPH and PAHs that would preclude reinjection of this water down-gradient of the dewatering area. An on site



treatment plant may be capable of reducing TPH and PAH concentrations in this water such that reinjection or disposal to sewer, as part of a trade waste agreement with Sydney Water, may be applicable. A pilot trial of possible treatment technologies was required to finalise the required approach;



A2 Scope of Works

In order to address the uncertainties listed in the previous Section, the following scope of work was completed at the site:

- Derivation of site specific assessment criteria for leachable concentrations that are considered suitable for the protection of environmental values;
- Assessment of the suitability of impacted material at the site to be remediated by stabilisation, through:
 - the excavation of testpits and sampling of impacted material. representative of that at the site;
 - o bench scale treatment of cement stabilisation on impacted material; and
 - analysis of treated material for leachable concentrations and unconfined compressive strength (UCS).
- Further characterisation of leachable contaminant concentrations through the collection and analysis of up to nine samples of impacted fill or soil from potential source areas;
- Assessment of the expected rate of dewatering required on site by completion of short interval pump tests at representative locations across the site; and
- Assessment of suitable disposal locations for water generated during dewatering and required treatment, if any, by:
 - Completion of a water treatment trial on water collected during the pump tests. Three treatment technologies are to be trialled, namely an oil/water separator, a sand filter and a granulated activated carbon (GAC) filter; and
 - Collection and analysis of samples of influent and effluent from each treatment method.

The methods and results of the pre-remediation testing program developed to address these data gaps are detailed in the subsequent sections.



A.3 Derivation of Site Specific Leachability Criteria

In preparing the revised remediation strategy for the site, it is proposed that both the total and leachable concentrations of the contaminants of concern are assessed in treated material prior to determining its suitability for on site reuse.

Determination of acceptable total concentrations in treated material is expected to be straightforward and based on human health related investigation levels for commercial land use.

Site specific leachability criteria were derived by the site specific criteria in the JBS letter 'Derivation of Site Specific Leachability Criteria – Former Macdonaldtown gasworks, Burren Street, Erskineville, NSW' Reference JBS 40913-1550, provided as **Appendix B.**

The site specific leachability criteria were derived based on the relevant criteria for protection of groundwater resources in the area, but also incorporating a dilution attenuation factor (DAF) to account for dilution of contaminated groundwater that occurs at the receiving water body, as it becomes mixed with groundwater discharged from the remainder of the catchment. A DAF value of 16.6 was determined for the site, and the resulting site specific leachability criteria for the site are provided in **Table A3.1** below.

Full details of the DAF and criteria derivation methods are provided in **Appendix B**.



Contaminant	Limit of Reporting	Aquatic Ecosystems ¹	Site Specific Criteria for assessment for leachable concentrations in soil ³
Arsenic (III/V)	0.1	2.3 ² / 4.5 ^{2, 5}	36.8
Cadmium	0.1	0.7 ¹	11.36
Chromium (III)	1	10	160
Chromium (VI)	0.1	4.4	70.4
Copper	0.1	1.3	20.8
Lead	0.1	4.4	70.4
Manganese	1	80 ²	1280
Mercury	0.05	0.1 ¹	1.76
Nickel	1	70	1120
Zinc	1	15	240
	,	VOLATILE ORGANIC COMP	OUNDS
Benzene		500	8000
Toluene		180 ²	2880
Ethylbenzene		5 ²	80
Xylene (M+O+P)		625 ²	10000
Styrene		1600	25600
Phenols		400	6400
	POLY	CYCLIC AROMATIC HYDRC	AOCARBONS
Benzo(a)Pyrene	0.1	0.1 ²	1.6
Naphthalene	0.1	70	1162
Phenanthrene	0.1	0.6 ²	9.6
Anthracene	0.1	0.01 ²	0.16
Fluoranthene	0.1	0.1 ²	1.6

Table A3.1: Site Specific Leachability Assessment Criteria (all units in μ g/L)

¹ 95% protection levels (marine ecosystems) have been used. When these levels fail to protect key test species, the 99% protection levels were used - ANZECC/ARMCANZ (2000). The 99% protection levels have been adopted in line with recommendations in Section 8.3.7 of ANZECC/ARMCANZ 2000.
 ² Insufficient data to derive a reliable trigger value. In these instances, reference has been made to low reliability trigger levels contained in ANZECC/ARMCANZ (2000).
 ³ Adopted criteria – 16x ANZECC/ARMCANZ 2000 trigger values, which is equivalent to 16.6x the LOR where anylicable

where applicable ⁴ Where no ANZECC/ARMCANZ 2000 trigger values, LOR used prior to applying dilution attenuation factor of 16



A.4 Sampling and Analysis Plan

Data Quality Objectives (DQOs) were developed for the remainder of the pre-remedial assessment works, as discussed in the following sections.

A4.1 State the Problem

RailCorp are committed to undertaking remediation works on the former Macdonaldtown Gasworks site. Contamination has been identified on the site consisting of free tar present in the abandoned gasworks infrastructure (tar wells, retort pipes etc.) and elevated concentrations of lead, B(a)P, Total PAH, TPH C₆-C₉, TPH C₁₀-C₃₆ and BTEX compounds in soils present from the surface to depths in excess of 10 m below ground level (bgl). Samples of groundwater collected from both the perched water table present in fill and from the bedrock aquifer underlying the site were reported to contain elevated concentrations of heavy metals, B(a)P, Total PAH, TPH C₆-C₉, TPH C₁₀-C₃₆ and BTEX. In accordance with the land use zoning, the objective of the proposed remediation program would be to render the site suitable for commercial land use with no unacceptable risk to the environment.

The pre-remedial assessment works were conducted to provide additional detail on the likely manner in which remediation is to be undertaken.

The team at JBS utilised for the project comprised:

- Andrew Lau Principal Contaminated Land, was the Project Director, responsible for client liaison and program design;
- Matthew Parkinson Principal Contaminated Land, provided specialist technical input to program design and assessment of results;
- Sumi Dorairaj Senior Environmental Consultant, was the Project Manager responsible for client liaison, site work planning and design, health and safety management, site work co-ordination, subcontractor management and reporting; and
- Tim Davis Senior Field Scientist, was responsible for completion of site works including subcontractor management, health and safety management and reporting.

The conceptual site model, prior to the completion of the pre-remedial assessment works, has been summarised in **Sections 2** of the main body of this report.

A4.2 Identify the Decision

The following decisions were made:

- For impacted material that remains *in-situ*, or that has been remediated through stabilisation, what are the maximum leachable concentrations of COC that present no additional risk to future users or to the surrounding environment via groundwater?
- For the coal tar impacted fill or soil to be remediated, what proportion (by mass) of calcium or magnesium based oxide cement must be added during treatment for immobilisation to be successful in reducing the leachable concentrations to acceptable levels?
- For the coal tar impacted fill or soil to be remediated, what proportion (by mass) of calcium or magnesium based oxide cement must be added during treatment to satisfy the requirements for disposal under the DECCW General Approval of the



Immobilisation of Contaminants in Waste 2005/14 (Coal Tar Wastes from Former Gas Works sites)?

- What are conservative and reasonable estimates of the dewatering rates that will be required on site during remediation of fill and/or soil?
- Out of an oil/water separator, sand filter and granular activated carbon (GAC) filter, which technology (or combination thereof) is best suited to treat the water generated on site during dewatering?

A4.3 Identify Inputs to the Decision

Inputs to the decisions were:

- A site conceptual model provided in the RAP(CH2M Hill, 2007);
- Existing groundwater data consisting of gauging, survey and limited analysis of groundwater samples;
- Soil environmental data collected by soil sampling and analysis;
- Stabilisation trial data comprising treated soil analysis and geotechnical strength testing for a range of treatment conditions;
- Groundwater pump test results from a representative number of wells on site consisting of gauging and pumping rate results to allow estimate rates of inflow into excavations;
- Treated groundwater environmental data generated by processing the water generated during the pump test through a trial scale water treatment plant, with sampling and analysis of recovered effluent;
- Soil criteria to be selected on the site based on a range of potential environmental values as defined by assessment criteria prepared in **Section A7.2**;
- Waste assessment criteria specified for Non-Liquid Waste in 'Waste Classification Guidelines, Part 1: Classifying Waste' (DECC, 2009) and also the 'General Approval of the Immobilisation of Contaminants in Waste' (NSW EPA 2005);
- Groundwater criteria selected on the site to be protective of current and potential future environmental values permitted under the site zoning, as defined by assessment criteria prepared in **Section A7.4**;
- Confirmation that data generated by sample analysis are of a sufficient quality to allow reliable comparison to assessment criteria as undertaken by assessment of quality assurance / quality control as per the data quality indicators established in Section 6.4; and
- Qualitative assessment of remediation methods commenced in Section A9.

A4.4 Define the Study Boundaries

The conditions to which the decisions applied and within which data were collected are summarised following:

- The lateral boundaries of the site are consistent with the boundaries of Part Lot 50 in Deposited Plan 1004167 that comprises the site;
- Risk goals as required to be met for on-going industrial use of the site and the future user populations that may be potentially impacted by contaminated material retained on the former Macdonaldtown gasworks site;



- Future site occupants and occupants in the proximity of the site, who have been assumed to be potentially exposed to groundwater present in the upper-most water bearing zone only; and
- Background groundwater quality was collected from the hydraulically upgradient boundary of the site, interpreted as being the western boundary of the former gasworks site.

A4.5 Develop a Decision Rule

The requirements for future remedial Strategy with respect to fill/soil stabilisation and groundwater treatment were proposed to be assessed by:

- Comparison of environmental data, from untreated and treated soil or fill, to criteria provided for the off site disposal of waste as available within 'General Approval of the Immobilisation of Contaminants in Waste' (NSW EPA 2005), Approval Number 2005/14 Coal Tar Waste from Former Gasworks Sites;
- Comparison of environmental data, from treated water and leachable concentrations in stabilised soil, to criteria provided for the protection of ecological ecosystems as available within:
 - 95% protection values for marine water ecosystems provided in ANZECC/ARMCANZ (2000) 'Australian and New Zealand Guidelines for Fresh and Marine Water Quality';
 - Dutch intervention levels provided for TPH to Netherlands Ministry of Housing, Spatial Planning and Environment (2000) 'Circular on target values and intervention values for soil remediation'; and
 - Site specific leachability criteria as provided in **Appendix B**;

as assessed at the location of the nearest potentially impacted marine ecosystems, *i.e.* Alexandra Canal.

A4.6 Specify Limits of Decision Error

There are two types of decision error identified in AS4482.1-2005 'Guide to the investigation and sampling of sites with potentially contaminated soil Part 1: Non-volatile and semi-volatile compounds'. These include:

- a) Deciding that the site is acceptable when it actually is not; and
- b) Deciding that the site is unacceptable when it is.

Limits are required to be set on each type of error presented here. AS4482.1-2005 nominates that a 5% probability of (a) and 20% probability of (b). These were adopted for this investigation.

To assess the usability of the data prior to making decisions, the data were assessed against pre-determined Data Quality Indicators (DQIs) for precision, accuracy, representativeness, completeness and comparability. The acceptable limit on decision error is 100% compliance with DQIs.

The pre-determined Data Quality Indicators (DQIs) established for the project are discussed below in relation to precision, accuracy, representativeness, comparability and completeness (PARCC parameters), and are shown in **Table A4.1**.

• **Precision** - measures the reproducibility of measurements under a given set of conditions. The precision of the laboratory data and sampling techniques is



assessed by calculating the Relative Percent Difference (RPD) of duplicate samples.

- **Accuracy** measures the bias in a measurement system. The accuracy of the laboratory data that are generated during this study is a measure of the closeness of the analytical results obtained by a method to the 'true' value. Accuracy is assessed by reference to the analytical results of laboratory control samples, laboratory spikes and analyses against reference standards.
- **Representativeness** –expresses the degree which sample data accurately and precisely represent a characteristic of a population or an environmental condition. Representativeness is achieved by collecting samples on a representative basis across the site, and by using an adequate number of sample locations to characterise the site to the required accuracy.
- **Comparability** expresses the confidence with which one data set can be compared with another. This is achieved through maintaining a level of consistency in techniques used to collect samples; ensuring analysing laboratories use consistent analysis techniques and reporting methods.
- **Completeness** is defined as the percentage of measurements made which are judged to be valid measurements. The completeness goal is set at there being sufficient valid data generated during the study.

Data Quality Indicators	Frequency	Data Quality Criteria
Precision		
Blind duplicates (intra laboratory)	1 / 20 samples	<30-50% RPD ¹
Blind duplicates (inter laboratory)	1 / 20 samples	<30-50% RPD ¹
Accuracy		
Surrogate spikes	All organic samples	60-140%
Laboratory control samples	1 per lab batch	60-140%
Matrix spikes	1 per lab batch	60-140%
Representativeness		
Sampling appropriate for media and analytes		-
Samples extracted and analysed within holding times.	-	pH (7 days), organics (14 days), inorganics (6 months)
Trip spike	1 per sampling event	70-130% recovery
Trip blank	1 per sampling event	<lor< td=""></lor<>
Comparability		
Standard operating procedures for sample collection & handling	All Samples	All samples
Standard analytical methods used for all analyses	All Samples	All samples
Consistent field conditions, sampling staff and laboratory analysis	All Samples	All samples
Limits of reporting appropriate and consistent	All Samples	All samples
Completeness		
Soil description and COCs completed and appropriate	All Samples	All samples
Appropriate documentation	All Samples	All samples
Satisfactory frequency and result for QC samples	All QA/QC samples	-
Data from critical samples is considered valid	-	Critical samples valid

Table A4.1 Summary of Quality Assurance / Quality Control Program

If any of the DQIs were not met, further assessment was necessary to determine whether the non-conformance significantly affected the usefulness of the data. Corrective actions include requesting further information from samplers and/or analytical laboratories, downgrading of the quality of the data or alternatively, collection of replacement data.



A4.7 Optimise the Design for Obtaining Data

The objectives of the works were to provide initial clarity on:

- Whether stabilisation will be a suitable remedial method to render the site suitable for commercial land use;
- Whether stabilised soil will be suitable for disposal off site in accordance with the requirements of IA 2005/14 (NSW EPA, 2005);
- whether the treated soil will be capable of being retained on site from both human health and environmental perspectives; and
- what allowances will need to be made for treatment of water generated during dewatering, in terms of the magnitude of water requiring treatment and type of treatment required.

All works were undertaken as per the methods specified in **Section A6**. The results of all additional work were assessed against historical results for contaminant concentrations in soil and slug test results in shallow wells. The results obtained are to be used to direct the design of the remediation programme for the site. Where the results of the trial works were inconclusive or inconsistent with the historical dataset, then to enable resolution, additional works have been included in the finalised remedial strategy for the site.



A.5 Sample Locations

As discussed in **Section A.2**, the pre-remedial trials were undertaken by a staged program of works. The program was staged to minimise the scale of subsurface disturbance to occur prior to remediation. The scope of works was as follows:

- Sampling of test pits excavated in the footprints of the Archaeological Research program pits:
 - In the vicinity of three former sampling locations, namely BHC, MG02 and MG10A, glass jar and bulk samples were collected at JBS TP3 4.0-4.2m, JBS TP3 1.3-1.7 m and JBS TP1 0.3-0.4 m depth respectively. Sampling depths were targeted to horizons previously demonstrated to be impacted by CoC. A duplicate bulk sample was collected at JBS TP3 4.0-4.2 m for analysis as a stabilisation trial control sample ; and
 - In the footprints of the remaining archaeological test pits, three samples were analysed from JBS TP2, JBS TP4 and JBS TP5;
- Completion of short interval pump tests at MW04S, MW06S, MW37S and MW42S
- Completion of a water treatment trial on water collected during the pump tests. Three treatment technologies are to be trialled, namely an oil/water separator, a sand filter and a GAC filter.

The methods to be employed in the collection and assessment of these data are detailed in **Section A.6.**



A.6 Assessment Methods - Soil Stabilisation Trials and Additional Leachability Testing

A6.1 Soil Sample Collection

JBS test pit locations are presented on **Figure A1**. All test pit locations were dug using a backhoe loader under the direction of trained JBS field staff. Soil/fill from the test pit above the target layer was stockpiled on plastic sheeting during excavation. Once the top of the target layer was identified, the material from this layer was stockpiled separately for sample collection. Proposed target layer depths were described in **Section A.5**, the actual sample collection depth was based on observed field conditions and are also summarised in **Section A.9**. Test pits were terminated at the base of the target layer or prior refusal.

For test pits where stabilisation testing was required, a 250 mL glass jar sample for analytical testing and a bulk sample (approximately 50kg) for treatment were collected from cuttings from the segregated target layer stockpile(s), placed in the appropriate sampling containers and sealed. Field duplicate jar samples and one duplicate bulk sample were also collected and analysed at the required frequency.

For test pits where samples were to be collected for analytical testing only, a 250 mL glass jar sample was collected directly from the excavator bucket, taking care to avoid collecting material in contact with the bucket walls.

A6.1.1 Soil Sample Containers

During the collection of soil samples, features such as seepage, discolouration, staining, odours and other indications of contamination were noted on field reporting sheets.

Test pit cuttings from the sampling layer were immediately transferred to sample containers of appropriate composition. For the bulk samples this comprised new, clean, plastic, bulk bags capable of containing up to 15 kg of soil without breach. For the analytical testing sample this comprised new, laboratory prepared, acid washed glass sample jars with Teflon lined screw closures. Labels on all sample containers were completed to record JBS job number; sample identification number; and date and time of sampling.

Filled, glass jar sample containers were sealed and transferred to a chilled ice box for preservation prior to and during shipment to the analytical laboratory. Bulk samples were maintained in cool, sheltered conditions at the JBS Office until delivery to the testing laboratory was possible. A chain-of-custody form was completed and forwarded with the samples to the laboratories. The following information was included on the chain–of-custody form:

- Sample identification;
- Signature of sampler;
- Date and time of collection;
- Place of collection;
- Type of sample;
- Number and type of container;
- Inclusive dates of possession; and
- Signature of receiver.



A6.1.2 Duplicate and Triplicate Sample Preparation

Field soil duplicate and triplicate samples were obtained during sampling using the following methods. Soil samples were divided into three samples with minimal disturbance to reduce the potential for loss of volatiles and placed in three laboratory prepared glass jars. All jars were filled with no headspace to reduce the potential for loss of volatiles and separately labelled as the primary, duplicate and triplicate samples before being placed in the same chilled ice box for transport to the laboratory.

A6.1.3 Decontamination of Soil Sampling Equipment

Soil sampling equipment used by JBS comprised only disposable gloves. The gloves were discarded following collection of each sample and a clean, new pair donned prior to handling the next sample material.

Test pits were excavated in order of the 'least likely contaminated' location to 'most likely contaminated' location to minimise the potential for cross contamination. At the completion and reinstatement of each testpit, the backhoe bucket was assessed for visible contamination and/or odours, and scrubbed and rinsed as appropriate.

A6.1.4 Stabilisation Testing Procedure

The glass jar sample of the untreated soil at each location was submitted to a NATA accredited laboratory for the analysis of total and leachable concentrations of heavy metals, B(a)P, cresol, BTEX, styrene and phenol.

A bench scale stabilisation trial was completed on the bulk sample collected at each test pit location. The samples were delivered to two laboratories for testing. Enviropacific Services Pty Ltd (Enviropacific) was subcontracted by JBS for completion of the stabilisation trial, and Cardno Bowler (Cardno) was subcontracted for completed of the clay content analysis, as follows:

- Preparation of a 20 kg subsample for clay content analysis by wet sieve analysis in accordance with AS 3686-1993 Test Sieve Analysis;
- Preparation of the remainder into 10 kg subsamples by removing the oversize components (*i.e.* passing the sample through at 20mm sieve) and noting descriptions of any oversize particles removed, foreign materials etc or other works required to complete the sieving;
- Subsamples were then divided and weighed out to form 5 kg samples per material. The subsampled material was sampled for laboratory analysis;
- Each 5 kg subsample was combined with calcium oxide based cement at the ratios of 5%, 12.5% and 20% (by mass) and mixed by hand for approximately 5 minutes;
- Water was added accordingly to achieve the moisture content required for the complete hydration of cement, noting the sample mass pre and post water addition;
- Samples were further mixed by hand for another 5 minutes;
- Mixed samples were cast into cylinders in accordance with AS 1012.8.1-2000 Methods of testing concrete – Method of making and curing concrete – Compression and indirect tensile test specimens;



- Cast cylinders were left to stand for approximately 24 hrs prior to wrapping and if required, transport to another NATA accredited laboratory, for curing in a fog room at 23°C for a further 6 days;
- All samples (*i.e.* 2 x 5 kg subsamples) shall be then tested in accordance with *AS1012.9-1999 Methods for testing concrete – Determination of the compressive strength of concrete specimens,* providing duplicate results for each treatment strength;
- Results of compressive strength analysis were compared to the requirements of NSW EPA General Immobilisation Approval 2005-14;
- A subsample of the treated material at each of the 5%, 12.5% and 20% cement strengths was collected and submitted to a NATA accredited laboratory for total and leachable concentrations of heavy metals, B(a)P, cresol, BTEX, styrene and phenol. ASLP testing was conducted on the treated soil samples to provide an indication of treated material performance under on site conditions; and
- Results of the analytical testing on treated and untreated samples were compared against the requirements of DECCW General Immobilization Approval 2005-14.

A6.2 Assessment Methods - Groundwater Pump Testing

The proposed pump test locations, MW03S, MW06S, MW07S and MW31 are shown on **Figure A1.** The proposed test methodology was a modified pumping test based on the procedure provided in MacDonald A, Barker J and Davies J (September 2008) '*The bailer test: a simple effective pumping test for assessing borehole success*' in <u>Hydrogeology</u> <u>Journal</u>. Data analysis was completed using the AquiferWin32 analysis package. The test was undertaken in the field by the following method:

- Install water level logger within monitoring well set up to record readings every second;
- Install groundwater pump to the lowest depth possible;
- Allow water level in well to stabilise for two minutes;
- Pump well for 10 minutes at maximum flow rate, or until pump is heard to go dry. Record time of pumping using stop watch;
- Maintain logger in position during well recovery, and remove once water level has recovered to 87% of pre-test level; and
- Remove pump and logger from well.

A 12V submersible electric pump was used for the tests. The maximum flow rates for pumps of this nature range from approximately 17 L.min⁻¹ at 1m depth to between 12 and 16 L.min⁻¹ at 6 m depth. The achieved flow rates, were however, dependent on the well and formation characteristics, and the stable maximum flow rates achieved are provided in **Section A.11.**

Given that the pump tests were targeted to the shallow perched aquifer, and the existing well network on the site is widely spaced and not systematic, the pumping well was used as the observation well for each of the tests in accordance with the modified pumping test method.

A6.3 Assessment Method - Groundwater Treatment Trial



Total Environmental Solutions Pty Ltd (TES) was subcontracted by JBS to construct a trial water treatment system (WTS) for site comprising an oil water separator, a sand filter and GAC filter. The WTS was constructed with taps in the pipe work connecting each treatment component to enable easy collection of water samples through the treatment process.

Samples of the trial water were collected at the influent and effluent points of each WTP component and submitted for analysis for TPH, PAHs, VOCs, Phenols and heavy metals. Measurements of water quality parameters, specifically Electrical Conductivity (EC), pH, temperature, dissolved oxygen (DO) and oxidation-reduction potential (ORP), were also collected for each water sample.

A6.3.1 Water Treatment Trial Sample Collection

The trial water was collected as grab samples for laboratory analysis. Sample bottles were filled directly from the influent and effluent taps installed on the system. A subsample was passed into a clean beaker for field filtering of samples for metals analysis. Bottles were filled to the top, and sealed tightly with the provided lids as soon as possible.

Immediately after sample collection, another subsample of the water was passed through a flow cell to allow for the measurement of water quality parameters.

During the collection of treatment trial samples, characteristics of the water such as colour, odour and apparent turbidity or any indications of contamination were noted on field reporting sheets. Phase separated hydrocarbons (PSH) were not observed during the groundwater pumping or treatment process.

Filled and sealed sample bottles were transferred to a chilled ice box for preservation prior to and during shipment to the analytical laboratory. A chain-of-custody form was completed and forwarded with the samples to the laboratories. The following information will be included on the chain–of-custody form:

- Sample identification;
- Signature of sampler;
- Date and time of collection;
- Place of collection;
- Type of sample;
- Number and type of container;
- Inclusive dates of possession; and
- Signature of receiver.

A6.3.2 Duplicate Sample Preparation

One duplicate sample was obtained during the field sampling event using the following methods. Two sets of sample bottles were filled with the sample water. Once full, samples were sealed with zero headspace to reduce the potential for loss of volatiles and separately labelled as the primary, duplicate and triplicate samples before being placed in the same chilled ice box for laboratory transport.

A6.3.3 Laboratory and Field Analyses

NATA accredited laboratories were used for all laboratory analysis and testing of soil and groundwater samples. The schedule of analysis undertaken on samples of soils and groundwater is summarised in **Table A6.1**.



Table A6.1:	Summary of Proposed Analysis Reg	Jime		
Media	Analysis Frequency	Analytes		
Soil	For each stabilisation treatment test pit sample (inc	cluding one control sample):		
	1 sample untreated soil or fill	Total and leachable (TCLP) heavy metals(As, Cd, Cr, Pb, Mn, Hg and Ni), PAHs, BTEX, cresol, styrene and phenol Clay Content by wet sieve analysis ²		
	subsample soil to be treated with 5% cement addition by mass	Total and leachable (ASLP ¹) heavy metals (As,		
	subsample soil to be treated with 12.5% cement addition by mass	Cd, Cr, Pb, Mn, Hg and Ni), PAHs, BTEX, cresol, styrene and phenol; and		
	subsample soil to be treated with 20% cement addition by mass	 Unconfined Compressive Strength (UCS) on cured sample² 		
	For each leachability sampling test pits location			
	Maximum of 3 samples per pit	Total and leachable (TCLP) heavy metals(As, Cd, Cr, Pb, Mn, Hg and Ni), PAHs, BTEX, cresol, styrene and phenol		
Water	For each water treatment technology tested:	-		
	1 sample of influent water; 1 sample of water resident in the system; and	Water quality parameters (EC, DO, pH, temperature and ORP) TPH		
	1 sample of effluent water.	PAHs (at 0.1 μ g.L ⁻¹) detection limits		
		VOCs		
		Phenols;		
		Heavy metals (As, Cd, Cr, Cu, Pb, Hg, Ni and Zn)		

Notes

1: ASLP testing to be undertaken using a sample of groundwater collected from an upgradient unimpacted well as the leaching media, test performed as per standard ASLP method.

2: Tests will be undertaken by a geotechnical laboratory subcontracted by JBS. Given the known contamination status of the material under assessment, any of the subcontractors' personnel who will come in contact with the samples will be inducted onto the JBS Job Risk Assessment for the works, and briefed regarding the necessary health and safety requirements.

A6.4 Laboratory Analyses

Envirolab Services Pty Ltd (Envirolab) at Chatswood, NSW was contracted as the primary laboratory for the required analyses. The secondary laboratory was SGS Australia Pty Ltd (SGS) at Alexandria, NSW. Both laboratories are NATA registered for the required analyses. In addition, the laboratories were required to meet JBS internal QA/QC requirements.

Geotechnical testing of bulk and treated samples was completed by Cardno and Douglas Partners Pty Ltd. Both laboratories are NATA accredited for the testing undertaken.



A7. Assessment Criteria

A7.1 Regulatory Guidelines

The investigation was undertaken with consideration to aspects of the following guidelines, as relevant:

- *Contaminated Sites: Guidelines for Assessing Service Station Sites,* NSW EPA, 1994 (EPA 1994)
- Contaminated Sites: Sampling Design Guidelines, NSW EPA, 1995 (EPA 1995)
- Contaminated Sites: Guidelines for Consultants Reporting on Contaminated Sites, NSW EPA, 1997 (EPA 1997)
- Contaminated Sites: Guidelines for the NSW Site Auditor Scheme, 2nd Edition, NSW EPA, 2006 (DEC 2006)
- National Environment Protection (Assessment of Site Contamination) Measure, National Environment Protection Council, 1999 (NEPC 1999)
- Australian and New Zealand Guidelines for the Assessment and Management of Contaminated Sites, Australian and New Zealand Environment and Conservation Council and the National Health and Medical Research Council, 1992 (ANZECC/NHMRC 1992)
- Australian and New Zealand Guidelines for Fresh and Marine Water Quality. Australian and New Zealand Environment and Conservation Council and Agriculture and Resource Management Council of Australia and New Zealand, Paper No 4, 2000 (ANZECC/ARMCANZ 2000)
- Australian Drinking Water Guidelines, National Health and Medical Research Council and Agriculture and Resource Management Council of Australia and New Zealand, 2004 (NHMRC/NRMMC 2004)
- Environmental Health Risk Assessment: Guidelines for assessing human health risks from environmental hazards, Department of Health and Ageing and EnHealth Council, Commonwealth of Australia, June 2002 (EnHealth 2002)
- Contaminated Sites: Guidelines for the Assessment and Management of Groundwater Contamination, NSW DEC, March 2007 (DEC 2007)
- Contaminated Sites: Guidelines on Duty to Report Contamination under the Contaminated Land Management Act 1997, NSW DECC, June 2009 (DECC 2009)



A7.2 Soil Criteria

Assessing the success of stabilisation in remediating soil is a two stage process, as follows:

- Results of geotechnical testing on treated material and results of total contaminant concentrations on untreated soil were assessed against the 'Conditions of Approval' as specified in 'General Approval of the Immobilisation of Contaminants in Waste' (NSW EPA 2005), Approval Number 2005/14 - Coal Tar Waste from former Gasworks Sites. The conditions are summarised in Table A7.1 and
- 2. Results of TCLP and ASLP testing on untreated and treated material were assessed against groundwater assessment criteria discussed in **Section 4.3**.

	Requirement (IA 2005/14, NSW EPA) ¹
Concentrations in untreated material	PAHs – 13 000 mg.kg ⁻¹ B(a)P – 500 mg.kg ⁻¹ Non-halogenated phenols – 2000 mg.kg ⁻¹ Total Cyanide 4000 mg.kg ⁻¹
Reagent used for stabilisation	Calcium or Magnesium Oxide 2:1
Maximum allowable stabilisation ratio	(<i>i.e.</i> by mass, 2 parts reagent to 1 part untreated soil)
UCS	1 MPa as the 95% lower confidence limit value (as assessed by AS1012.9-1999)

Table A7.1:	Soil Criteria (units as specified)	
	Son cinceria (units as specifica)	

Additionally, the results of clay content testing on the untreated material were assess to determine whether this property is likely to affect the efficacy of stabilisation as the remediation method.

Should the treated material fail to comply with the requirements of **Table A7.1**, the material will need to be assessed against the criteria provided for off site disposal of waste as available within *Waste Classification Guidelines*, *Part 1: Classifying Waste'* (DECC, 2009);

A7.3 Soil Leachability Data

Soil leachability data was assessed against the :

- the site specific leachability criteria specified in Table A3.1; and
- criteria provided for off site disposal of waste as available within *Waste Classification Guidelines, Part 1: Classifying Waste'* (DECC, 2009).

A7.4 Treated Groundwater Criteria

Treated groundwater data were assessed against groundwater investigation levels (GILs). DEC (2007) '*Guidelines for the Assessment and Management of Groundwater Contamination*' instructs that GILs adopted for a site are required to take into consideration groundwater environmental values.

Environmental values are defined in ANZECC / ARMCANZ (2000) as '...particular values or uses of the environment that are important for a healthy ecosystem or for public benefit, welfare, safety or health which require protection from the effects of pollution, waste discharges and deposits'.

The site lies within the lower estuaries of the Sydney Harbour and Parramatta River catchment boundary established by the DECC



(<u>http://www.environment.nsw.gov.au/ieo/SydneyHarbour/map.htm,</u> as on 17 June 2010). The interim water quality objectives established in NSW Government (2006) '*Environmental Objectives for Water Quality and River Flow* for waterways affected by urban development in the Sydney Harbour and Parramatta River catchment include:

- Protection of aquatic ecosystems;
- Protection of visual amenity;
- Protection of secondary contact recreation; and
- Protection of primary contact recreation in the long term, *i.e.* in 10 years or more.

These water quality objectives are required to be applied at the point of ecological discharge, which previous investigations have stated would include Alexandra Canal located to the south-east of the site.

A discussion of the use of groundwater in urban areas for drinking or agricultural purposes is provided in Sydney Coastal Councils Group (September 2006) 'Groundwater Management Handbook' (SCCG, 2006). It is stated there that "DNR advises that groundwater from individual domestic bores or spearpoints in urban areas <u>should never</u> be considered suitable as a drinking water supply. The health risk posed by untreated water from these installations can be significant. DNR also recommends that groundwater should not be used for watering edible produce (i.e. vegetables and fruits), nor for filling swimming pools and spas in an urban area. Any of these proposed uses requires frequent rigorous testing and treatment of groundwater to confirm its suitability for the intended purpose".

Drinking water will be considered as a potential future use in the event that economical yields from the bedrock may one day be achievable. However, noting the commentary provided in SCCG (2006) it is considered inappropriate to strictly assess concentrations of contaminants in groundwater collected from the Site against drinking water criteria. Similarly, strict assessment of concentrations of contaminants in groundwater collected from the site against primary and secondary contact recreation is inappropriate.

The following criteria were considered to allow each of these beneficial uses to be assessed:

- 95% protection levels for marine waters provided in ANZECC/ARMCANZ (2000) to be assessed at the nearest down gradient receptor;
- Recreational water quality criteria provided in Section 5.2.3 of ANZECC/ARMCANZ (2000) to be assessed at the nearest down gradient receptor;
- Recreational and aesthetic water quality criteria provided in Section 5.2.3.3 of ANZECC/ARMCANZ (2000) to be assessed at the nearest down gradient receptor;
- Irrigation water quality criteria provided in Section 9.2.5 of ANZECC / ARMCANZ (2000) to be applied as a long term objective on the site boundary; and
- Drinking water criteria provided in NHMRC NRMMC (2004) 'Australian Drinking Water Guidelines' to be applied as a long term objective at the site boundary and in consideration of the discussion provided earlier.

It is noted that the above summarised Australian guidelines do not provide guidelines for several substances. In these cases, DEC (2007) instructs that laboratory detection limits should be used as preliminary investigation levels. For substances with no drinking water criteria, reference may also be made to 'tap water criteria' provided to US EPA Region 9 (2004) 'Preliminary Remediation Goals'. It is noted that these are intended as screening level (preliminary) criteria, and may be based on toxicological data and exposure



parameters that may be less conservative than those which may be endorsed for use in Australia. The criteria used for assessment of soil leachability and groundwater data are summarised in **Table A7.2**.



	Aquatic Ecosystems ¹	Recreation ³	Visual Amenity⁴	Drinking Water ⁹	Irrigation ¹⁰	LOR ⁷
Total Petroleum Hydrocar	bons					
ГРН (C ₆ – C ₃₆)	7 ²	-	No sheen or odour	90 ¹¹	-	250 ⁷
Monocyclic Aromatic Hydr	ocarbons		<u>.</u>			
Benzene	500	10	-	1	-	
Toluene	180 ²	-	-	800	-	
Ethylbenzene	5 ²	-	-	300	-	
Xylene (M+O+P)	625 ²	-	-	600	-	
Isopropylbenzene	30 ²	-	-	660 ⁶	-	
n-propylbenzene	-	-	-	240 ⁶	-	
1,3,5-trimethylbenzene	-	-	-	12 ⁶	-	
1,2,4-trimethylbenzene	-	-	-	12 ⁶	-	
Tert-butylbenzene	-	-	-	240 ⁶	-	
p-isopropyltoluene	-	-	-	-	-	1
n-butylbenzene	-	-	-	240 ⁶	-	
Trihalomethanes	•				• 	
Bromodichloromethane	-	-	-	250	-	
Halogenated Aromatic Co	mpounds					
1,2-Dichlorobenzene	160	-	-	1,500	-	
1,4-Dichloeobenzene	60	-	-	40	_	
Halogenated Aliphatic Cor	npounds					
Vinyl chloride	100	_	_	0.3	_	1
1,1-Dichloroethane	250 ²	-	-	810 ⁶	_	
1,1-Dichloroethene	700 ²	-	_	30	_	
Cis-1,2-dichloroethene	700 ²	_	_	60	_	
Trans-1,2-dichloroethene	700 ²	_	_		_	
1,1,1-Trichloroethane	270 ²	_	_	3200 ⁶	_	
Trichloroethene	330 ²	30	_	0.0286	_	1
Tetrachloroethene	70 ²	10		50	_	
1,2,3-trichloropropane	-	-	_	0.0056 ⁶	_	1
Heavy Metals		-	-	0.0050		1
Arsenic (III/V)	2.3 ² / 4.5 ^{2, 5}	50	_	7	100	
Cadmium	0.7 ¹	50	-	2	100	
Caamium Chromium (III)	10	5	-	2	-	
	4.4	50	-	50	- 100	
Chromium (VI)		1000	-	50		
Copper	1.3	1000	-	2,000	200	
Lead	4.4 80 ²	50	-	10 500	2000	
Manganese		100	-	500	200	
Mercury	0.1 ¹	1	-	1	2	
Nickel	70	100	-	20 2000 ⁸	200	
Zinc	15	5000	-	3000 ⁸	2000	
Polycyclic Aromatic Hydro	i					
Naphthalene	50	-	-	6.2 ⁶	-	
Acenaphthylene	-	-	-	-	-	0.1
Acenapthene	-	-	-	370 ⁶	-	
Fluorene	-	-	-	240 ⁶	-	

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	Aquatic Ecosystems ¹	Recreation ³	Visual Amenity⁴	Drinking Water ⁹	Irrigation ¹⁰	LOR ⁷
Phenanthrene	0.6	-	-	-	-	0.1
Anthracene	0.01 ²	-	-	1800 ⁶	-	
Fluoranthene	1 ²	-	-	1500 ⁶	-	
Pyrene	-	-	-	180 ⁶	-	
Benzo(a)anthracene	-	-	-	0.092 ⁶	-	
Chrysene	-	-	-	9.2 ⁶	-	
Benzo(b&k)fluoranthene	-	-	-	0.092 ⁶	-	
Indeno(1,2,3-cd)perylene	-	-	-	0.092 ⁶	-	
Benzo(g,h,i)perylene	-	-	-	-	-	0.1
Dibenz(a,h)anthracene	-	-	-	-	-	0.1
Benzo(a)pyrene	0.1 ²	0.01	-	0.01	-	0.1

1. 95% protection levels (marine ecosystems) have been used. When these levels fail to protect key test species, the 99% protection levels were used - ANZECC/ARMCANZ (2000).

2. Insufficient data to derive a reliable trigger value. In these instances, reference has been made to low reliability trigger levels contained in ANZECC/ARMCANZ (2000).

3. Recreational purposes - Table 5.2.3 ANZECC/ARMCANZ (2000)

4. Recreational water quality and aesthetics – s.5.2.3.3 ANZECC/ARMCANZ (2000)

5. Criteria for As (V) selected

6. US EPA Region 9 (2004) tap water criteria

7. Laboratory limit of reporting provided for substances with insufficient published ecological / health investigation guidelines, or where published guidelines fall below laboratory limit of detection.

8. Based on aesthetic considerations. No Health based guideline published.

9. NHMRC NRMMC (2004) 'Australian Drinking Water Guidelines'

10. ANZECC / ARMCANZ (2000) Section 9.2.5., long term vales used.

11. Lowest of fraction specific criteria provided to WHO (2005) 'Petroleum Products in Drinking Water'



A.8 Pre-Remedial Investigations - Quality Assurance / Quality Control

A8.1 Soil QA/QC Results

The QA/QC results for soil samples collected at the site are summarised in **Table A8.1** and discussed in **Section A8.2** below. Detailed laboratory QA/QC results are included in the laboratory reports in **Appendix F** of the Revised Remedial Strategy document. Tabulated QA/QC results are provided in **Appendix G** of the Revised Remedial Strategy document.

Data Quality Indicator	Frequency	Results	DQI met?
Precision			
Soil Blind duplicates (intra laboratory)	1 field duplicate analysed – 9% of total samples >5% acceptable	0 - 121% RPD	Partial ¹
Soil Blind replicates (inter laboratory)	1 blind replicates analysed – 9% of total samples >5% acceptable	0-159% RPD	Partial ¹
Laboratory Duplicates	1 laboratory duplicate analysed depending upon analyte – 9% of total samples >5% acceptable	0-29% RPD	yes
Trip spike	1/1 batches (<lor)< td=""><td>96-100%</td><td>Yes</td></lor)<>	96-100%	Yes
Trip blank	1/1 batches (<lor)< td=""><td><lor< td=""><td>Yes</td></lor<></td></lor)<>	<lor< td=""><td>Yes</td></lor<>	Yes
Accuracy			
Surrogate spikes	All organic samples	39-130% recovery	Partial ¹
Matrix spikes	2/11 – 18% >5% acceptable	42-130% recovery	Partial ¹
Laboratory control samples	2/11 – 18% >5% acceptable	60-130% recovery	Yes
Representativeness			
Sampling appropriate for media and analytes	All samples	All sampling conducted in accordance with JBS procedures	Yes
Laboratory blanks	1/batch	<lor< td=""><td>Yes</td></lor<>	Yes
Samples extracted and analysed within holding times.	AII	All samples were extracted and analysed within holding times for the target analytes.	Yes
Comparability			
Standard operating procedures used for sample collection & handling	All	A team of three field staff used same standard operating procedures throughout works	Yes
Standard analytical methods used	All	Standard analytical methods used	Yes
Consistent field conditions, sampling staff and laboratory analysis	All	Sampling was conducted by one field staff using standard operating procedures in the same conditions throughout the works. The primary lab and secondary labs remained consistent throughout the investigation.	Yes
Limits of reporting appropriate and consistent	All	Limits of reporting were consistent and appropriate.	Yes
Completeness			
Soil description & COCs completed	All	All bore logs and COCs were completed appropriately.	Yes
Appropriate documentation	All	All appropriate field documentation is included in the Appendices.	Yes
Satisfactory frequency/result for QC samples	All	The QC results are considered adequate for the purposes of the investigation.	Yes
Data from critical samples is considered valid	All	Data from critical samples are considered valid.	Yes

Table A8.1 - Soil QA/QC Results Summary

¹ See discussion of DQI outliers below.



A8.2 Soil QA/QC Discussion

A8.2.1 Precision

With the following exceptions, all field duplicates recorded relative percentage differences (RPDs) within the acceptable range of less than 30-50%:

- In field replicate pair JBS TP1/0.3-0.4 m and QC2, Cd of 1.1 and 2.1 mg/kg respectively (RPD, 63%);
- in field duplicate pair JBS TP1/0.3-0.4 m and QA2 all individual PAH compounds with the exception fluorene, had RPDs in the range of 80 and 129 %;
- in field replicate pair JBS TP1/0.3-0.4 m and QC2 all individual PAH compounds with the exception acenaphthene, reported RPDs in the range of 51 and 159 %

The elevated RPDs appear to be due to the heterogeneity of the fill material sampled at these locations. The elevated RPDs are considered not to significantly affect the reliability of the data set because the concentrations reported for these analytes in the duplicate pairs were either both significantly higher or both significantly lower than the assessment criteria. In no field duplicate pair exceeding allowable RPDs, did the assessment criteria value lie between the two reported concentrations. Therefore the difference in concentrations in these field duplicate pairs did not affect the assessment of site suitability at these locations. As a further level of conservatism, only the higher of the two values reported in each field duplicate pair was used in the assessment dataset.

A8.2.2 Accuracy

With the exception of the following, all surrogate spike recoveries reported were all within the acceptable range:

- Phenol-d6: 55% recovery in JBS TP2/0.4-0.5 m and 57% recovery in JBS TP2/1.4-1.5 m;
- 2-fluorophenol: 59% recovery in JBS TP1/0.3-0.4 m and 57% recovery in JBS TP2/1.4-1.5 m;
- Phenol-d6: 55% recovery in JBS TP2/0.4-0.5 m and 57% recovery in JBS TP2/1.4-1.5 m; and
- 2,4,6-Tribromophenol: 59% recovery in JBS TP1/0.3-0.4 m, 39% in JBTP4/1.6 1.7 m and 55% recovery in JBS TP4/1.0 m

With the exception of 34% and 42% recoveries for 4-nitrophenol, all matrix spike recoveries reported were all within the acceptable range.

These spike exceedances are considered not to significantly affect the accuracy of the dataset given that phenol concentrations were below the laboratory detection limit in all field samples analysed.



A8.3 Groundwater Treatment Trial QA/QC Results

The QA/QC results for groundwater samples collected at the site are summarised in **Table A8.2** and discussed in **Section A8.4** below. Detailed laboratory QA/QC results are included in the laboratory reports in **Appendix F**.

Data Quality Indicator	Frequency	Results	DQI met?	
Precision				
Groundwater Blind duplicates (intra laboratory)	0 field duplicates analysed -0% of total samples >5% acceptable	-	No ¹	
Groundwater Blind replicates (inter laboratory)	1 blind replicates analysed – 9% of total samples >5% acceptable	0-13% RPD	Yes	
Laboratory Duplicates	No batch specific duplicates analysed	-	Yes	
Trip spike	1/1 batch (<lor)< td=""><td>95-129%</td><td>Yes</td></lor)<>	95-129%	Yes	
Trip blank	1/1 batch (<lor)< td=""><td><lor< td=""><td>Yes</td></lor<></td></lor)<>	<lor< td=""><td>Yes</td></lor<>	Yes	
Accuracy				
Surrogate spikes	All organic samples	60-140% recovery	Yes	
Matrix spikes	1 sample - acceptable	95-120%	No ¹	
Laboratory control samples	2 analysed – acceptable	70-130% recovery	Yes	
Representativeness				
Sampling appropriate for media and analytes	All samples	All sampling conducted in accordance with JBS procedures	Yes	
Laboratory blanks	1/batch	<lor< td=""><td>Yes</td></lor<>	Yes	
Samples extracted and analysed within holding times.	All	All samples were extracted and analysed within holding times for the target analytes.	Yes	
Comparability				
Standard operating procedures used for sample collection & handling	All	One staff member used same standard operating procedures throughout works	Yes	
Standard analytical methods used	All	Standard analytical methods used	Yes	
Consistent field conditions, sampling staff and laboratory analysis	All	Sampling was conducted by one staff member using standard operating procedures in the same conditions throughout the works. The primary lab and secondary labs remained consistent throughout the investigation.	Yes	
Limits of reporting appropriate and consistent	All	Limits of reporting were consistent and appropriate.	Yes	
Completeness				
Sample description & COCs completed	All	All field data sheets and COCs were completed appropriately.	Yes	
Appropriate documentation	All	All appropriate field documentation is included in the Appendices.	Yes	
Satisfactory frequency/result for QC samples	All	The QC results are considered adequate for the purposes of the investigation.	Yes	
Data from critical samples is considered valid	All	Data from critical samples are considered valid.	Yes	

Table 8.2 – Groundwater QA/QC Results Summary

¹ See discussion of DQI outliers below.

A8.4 Groundwater Treatment Trial QA/QC Discussion

A8.4.1 Precision

The field replicates recorded relative percentage differences (RPDs) within the acceptable range of less than 30 to 50%. It is noted that no field duplicate sample was analysed a part of the assessment, due to a shortage of sample containers on the day of the water treatment trial. Given that the data collected was used to assess treatment technologies,



and is not intended to characterise the site the lack of a field duplicate is considered not to affect the precision of the treated waste data set.

A8.4.2 Accuracy

All surrogate recovery results and matrix spike results were within the acceptable range.

A8.4.3 QA/QC Conclusion

The field sampling and handling procedures produced QA/QC results which indicated that the soil and water data are of an acceptable quality and suitable for use in site characterisation.

The NATA certified laboratory Certificates of Analysis indicated that the project laboratory was generally achieving levels of performance within its recommended control limits during the period when the samples from this program were analysed.

On the basis of the results of the field and laboratory QA/QC program, the soil and groundwater data are assessed to be of an acceptable quality upon which to draw conclusions regarding the environmental condition of the site, within the limitations of this study.



A.9 Pre-Remedial Assessment Results

Results of laboratory analysis on samples collected from the site are discussed in the following section. Sample location identifiers comprise the letters 'TP' (test pit) or 'MW' (monitoring well) as a prefix and a number as a suffix. A second suffix of 'S' or 'D' has been assigned to samples collected from shallow or deep wells respectively.

A9.1 Deviations from the Sampling, Analysis, Quality Plan

A Sampling, Analysis, Quality Plan (SAQP) was prepared for the Pre-Remedial Investigations documenting the proposed strategy and methods. During completion of the site works certain changes were made based on observed conditions on site and are discussed below.

<u>Changes to depth of bulk sample collect for cement stabilisation trial.</u> The proposed sampling depths listed in the SAQP were based on layers of fill/soil reported to contain highly elevated contaminant concentrations in previous assessments of the site. The actual sampling depth required in the current works was assessed by comparing the previous bore logs with the field observed material until a close match was apparent. **Table A9.1** summarises the proposed and actual bulk sample collection depths.

Proposed loca	tion	Actual Location	
Former Location ID / sample depth	Layer description (CH2M Hill, 2007)	New testpit ID / sample depth	Layer description (JBS, 2010)
MG10A/0.7m	Fill – black ash, and coke gravel	JBS TP1/0.3-0.4m	Fill- silty gravelly sand comprising coke, ash and slag, dark brown to black, dry to damp, some ballast gravel and cobble inclusions (irregular, basalt, hard)
MG02/ 1.8m	Fill – silty clay (original surface (?), spongy, wet, dark brown to black, low plasticity, black ooze, tar	JBS TP3/1.3-1.7m	Fill – silty clay dark brown, low plasticity, wet, heterogeneous with coke gravel and black tar ooze inclusions, strong PAH odours and black sheen on material
BHC (angled borehole)/ 6.0m	Fill – mixture of ironstone gravels and clays, black stains, wet sloppy, pierced through brick base annulus, free tar in bricks, saturated soil	JBS TP3/4.0 – 4.2m	Fill – silty clay yellow red with grey mottles, medium plasticity, very strong, wet PAHs odours, black ooze seepage throughout

Table A9.1: Proposed and Actual Bulk Sample Collection Depths

It is noted that while co-ordinates of the original sampling locations were used to generally locate the new test pit positions, heritage restrictions prevented placement of JBS TP3 close enough to the northern gasholder such that material at the base of the brick annulus could be sampled (as described in the original BHC bore log). The material sampled at depth in JBS TP3 was, however, considered to be consistent with tar impacted material likely to have originated from material used to fill the northern gasholder.

<u>Changes to groundwater wells to be used for pump tests and water collection for the treatment trial.</u> The SAQP nominated that wells MW07S, MW06S, MW03S and MW31 be used for pump tests and to collect groundwater for the water treatment trial. On commencement of site works the following was noted in relation to these wells:

- MW03S was unable to be located. Site plan indicated the well was placed in the vicinity of several large soil stockpiles, and it is uncertain whether this well remains viable;
- MW06S ran dry after extraction of approximately 40 L on both 7 and 9 July 2010;
- MW07S ran dry after extraction of approximately 50 L on both 7 and 9 July 2010; and



• MW31 was unable to be located at the time of the trial and it is likely this well has been destroyed.

Given the limited time available for completion of the trial, the majority of water for the trial was pumped from MW04S and MW37S. A total of 4000 L was pumped over the trial period from these two wells.

A9.2 Field Observations

Visual and/or olfactory indicators of contamination were noted at all sampling locations on the site. At the test pit locations:

- Ash, coke and slag inclusions were observed in JBS TP1 in shallow fill between 0 and 0.6 m depth;
- Ash, coke and slag inclusions were observed in JBS TP2 in shallow fill between 0 and 0.9 m depth. Coal tar odours were noted in all material encountered at this location to the termination depth of 1.8m, and a black oily sheen/ooze was observed on material between 1.3 and 1.8 m depth;
- Coke gravels, coal tar odours and a black sheen were noted on materials JBS TP3 from 1.3 m depth to the test pit termination depth of 4.3 m;
- Ash, slag and potential ACM inclusions were observed in JBS TP4 in shallow fill between 0 and 0.6 m depth. Coal tar odours were noted in fill between 0.9 and 1.4 m depth at this location; and
- Ash, coke and slag inclusions were observed in JBS TP5 in shallow fill between 0 and 0.5 m depth. Below this depth , coal tar odours were noted in all material encountered to the termination depth of 2.1 m.

In the shallow wells pumped for the water treatment trial, groundwater on the site was noted to be black grey in colour, highly turbid and having strong coal tar odours.

Test pit logs are included in **Appendix H**. Field notes from the pump tests are provided in **Appendix I**.

A9.3 Stratigraphy

The profile encountered across the site comprised fill overlying natural clay soils. The profile encountered is summarised in **Table A9.2** below.

Layer	Locations	Description	Depth Encountered (m bgl)
Fill	All boreholes	Dark brown, grey and black, gravelly silty ash, coke and slag inclusions, some materials	0 – 4.3
Free Groundwater	JBS TP2	0.9-1.0 m seepage entering pit From 1.0m onward material wet	0.9-1.8m ⁺
	JBS TP3	Materials wet from 0.4m onwards	0.4—4.3m ⁺

Table A9.2 – Summary of subsurface profile encountered during testpitting

Notes: + wet material extended beyond test pit termination depth.

The fill encountered in all test pits excavated on the site comprised layers of heterogeneous silty sand and gravelly silty sand, generally containing ash, slag and coke inclusions. Fill present in the top metre appeared to be highly heterogeneous with the majority of this top layer comprising anthropogenic materials. Generally with depth the portion of inclusions present in fill appeared to decrease, while moisture content and the intensity of coal tar odours appeared to increase. Black sheen or ooze was only encountered in fill material at depth.



A9.4 Soil Analytical Results

The soil sampling locations are shown on **Figure A.1** and summarised laboratory results are presented in **Tables A9.3 to A9.10**. Detailed analytical laboratory reports and chain of custody documentation are provided in **Appendix E**. Results of the stabilisation trial are presented in **Appendix I**. Results of clay content analysis are presented in **Appendix J**.

Laboratory results are discussed in the following sections.

A9.4.1 Concentrations in Soil in Untreated Material

Total and leachable concentrations in selected untreated soil samples are provided in **Tables A9.3 to A9.5**. Only detected contaminants have been tabulated, with full results for all undetected compounds contained in the laboratory certificate.

Despite having similar material descriptions, the concentrations contained in the samples analysed as part of the current program (JBS TP1 and JBS TP3) were less than the concentrations detected in the original test locations samples (BHC, MG10A and MG02 from CH2M Hill 2000). While part of this difference may be attributable to variations in the contaminant profile across the site, the major factor is considered likely to be a result of the difference in testing methods adopted. Specifically that the original samples were collected from drilled boreholes, while the current samples were collected from test pits. As the original samples appear to have been collected from thin, highly impacted layers it is likely that these concentrations over state the requirements of remediation. During remediation it is likely that these thin highly impacted layers will undergo some dilution/homogenisation while being excavated and stockpiled with less impacted material. The current samples, collected from test pits cuttings, are therefore considered more likely to represent the condition of free tar impacted materials during bulk excavation.

All heavy metals concentrations were less than the remediation acceptance criteria to be applied to the site. Phenols were also below detection limit in all samples and asbestos was not identified in any soil samples.

Concentrations of Total PAHs, B(a)P and /or naphthalene exceeded the remediation acceptance criteria in samples collected from JBS TP2/0.4-0.5 m, JBS TP2/1.4-1.5 m, JBS TP3/1.7 m, JBS TP4/1.6-1.7 m and JBS TP5/0.5 m.

In general the current results indicate that B(a)P concentrations were highest at shallow depths. The highest concentration of B(a)P was detected in sample JBS TP2/0.4-0.5 m comprising shallow fill with coke and slag inclusions, containing a total concentration of B(a)P of 64 mg/kg. Despite the high total concentration the leachable B(a)P concentration in this sample was below the laboratory detection limit. Furthermore, as the leachable concentrations of B(a)P all samples were below the laboratory detection limit, the data suggests that some degree of natural immobilisation is occurring. Review of historical site data presented in the RAP (CH2M Hill, 2007) indicates that only 10 other samples collected from the site contained B(a)P concentrations in excess of 64 mg/kg.

The most prominent feature of the detected PAH concentrations, are the relatively high naphthalene values, which are consistent with by-products resulting from the generation of gas from coal. Highest detected concentrations were generally associated with material observed to contain 'black ooze' and/or have a sheen present, and the elevated naphthalene concentrations were considered the source of the strong 'coal tar' odours noted during sampling. While the current dataset is limited, the results suggest the prevalence of naphthalene in soils may be linked to former use of the site. Naphthalene accounted for more than half of the total PAH concentration in samples JBS TP2/1.4-1.5 m and JBS TP3/1.7 m, collected from the anticipated tar source area in the vicinity of the



Northern Gasholder and retort area. In samples collected from the areas external to the Northern Gasholder and Retort (from TP1, TP4 and TP5), naphthalene accounted for a much lower portion of the Total PAH concentration. This appears consistent with PAH results for the site provided in Tables 1 to 8 of '*Delineation & Characterisation Sampling and review of Remedial Options, Former Macdonaldtown Gasworks - Burren Street, Erskineville*' (CH2M Hill, March 2007). Results in Tables 1 and 2 of that document (samples from the gasholder and retort area) indicate that where PAHs were detected, naphthalene was detected at much higher concentrations than any of the other PAH compounds. Results in Tables 3 to 8 of that document, providing results from the other areas of the site, show that naphthalene concentrations in these areas account for a small percentage of the Total PAH concentrations.

Additionally, it is noted that while the elevated total concentrations of naphthalene that exceeded the remediation were detected in samples JBS TP2/0.4-0.5 m, JBS TP2/1.4-1.5 m, JBS TP3/1.7 m, JBS TP4/1.6-1.7 m and JBS TP5/0.5 m, the corresponding leachable concentrations were all less than the JBS site specific leachability criteria with the exception of 2.8 mg/L in sample JBS TP2/1.4-1.5 m and 3.6 mg/L in sample JBS TP3/1.7 m.

BTEX compounds may also be generated during gas production from coal, however only low BTEX compounds were detected in the samples analysed from the site. Only one sample JBS TP2/1.4-1.5m contained a concentration above the assessment criteria, a total xylene concentration of 66 mg/kg. It is noted that the corresponding leachable concentration in this sample was below the laboratory detection limit.

A9.4.2 Conclusions Relating to Treatment Suitability

In samples exceeding the assessment criteria for B(a)P, naphthalene and/or total PAHs, the concentrations were less that the allowable limits for cement stabilisation provided in IA 2005/14. This suggests that the material present at JBS TP2/0.4-0.5 m, JBS TP2/1.4-1.5 m, JBS TP3/1.7 m, JBS TP4/1.6-1.7 m and JBS TP5/0.5 m are suitable for treatment by stabilisation and, once treated, suitable to be assessed under IA 2005/14.

The presence of total benzo(a)pyrene above the site assessment criteria in samples JBS TP2/0.4-0.5 m, JBS TP2/1.4-1.5 m, JBS TP4/1.6-1.7 m and JBS TP5/0.5 m, suggests that bioremediation is unlikely to effectively treat these materials for on site reuse. Bioremediation may however be conducted on these materials such that a reduced waste classification is achieved for off-site disposal.

The presence of elevated total xylenes in the sample JBS TP2/1.4-1.5 m is suitable for treatment by either cement stabilisation or bioremediation.

A9.4.2 Soil Treatment Trial

Following sample collection, material were delivered to Enviropacific for treatment by cement stabilisation. On receipt, the materials were processed, *i.e.* particles over 20 mm in diameter removed, with the remaining sample homogenised, subdivided and then treated with either 5, 12.5 or 20 % cement. At the request of the site Auditor, subsamples of each material were analysed after processing and subdivision, prior to the addition of cement. Samples were also analysed at the completion of the treatment trial. Contaminant concentrations in the soil through the treatment process are presented in **Appendix J.**

Mean values for each of the treated material samples are summarised in **Tables A9.6 and A9.7**. Full results are included in **Appendix J**.





Table A9.3 PAH, Phenol and Asbestos Concentrations in Untreated Soil Samples (units as specified)										
Sample ID (summarised material description)	Total B(a)P (mg/kg)	Leachable B(a)P (mg/L)	Total Naphthalene (mg/kg)	Leachable naphthalene (mg/L)	Total PAHs (mg/kg)	Leachable PAHs (mg/L)	Total Phenols (mg/kg)	Total Phenols (mg/L)	Asbestos	
JBS TP1/ 0.3-0.4m ¹ Fill sandy gravel some ash, coal and slag (Material 1)	3	<0.001	0.6	<0.001	30.1	BDL	BDL	BDL	Not identified	
JBS TP2/ 0.4-0.5m Fill – silty gravelly sand, coal tar odour, coke and slag inclusions	64	<0.001	13	0.022	770.2	0.037	BDL	BDL	Not identified	
JBS TP2/1.4-1.5m Fill – silty clay, with black tar ooze inclusions coal tar odour, coke and slag inclusions	7.7	<0.001	350	2.8	467.7	2.9	BDL	BDL	Not identified	
JBS TP3/1.7m ¹ Fill - silty clay dark brown, s with coke gravel and black tar ooze inclusions, strong PAH odours and black sheen on material (Material 2)	0.6	<0.001	310	3.6	330.2	3.742	BDL	BDL	Not identified	
JBS TP3/4.0-4.2m ¹ Fill - very strong PAHs odours, wet, black ooze seepage throughout (Material 3)	4.5	<0.001	1.7	0.066	30.6	0.089	BDL	BDL	Not identified	
JBS TP4/0.5m Fill – silty sand, ash slag and glass fragments, suspected ACM fragment	4.4	<0.001	<0.1	0.002	33.2	0.002	BDL	BDL	Not identified	
JBS TP4/1.0m Fill – silty clay slight coal tar odour	<0.05	<0.001	<0.1	<0.001	BDL	BDL	BDL	BDL	Not identifie	
JBS TP4/1.6-1.7m Fill – silty clay	30	<0.001	6.3	<0.001	349.8	0.003	BDL	BDL	Not identified	
JBS TP5/0.5m Fill – silty clay, with strong coal tar odours	30	<0.001	250	0.64	724.3	0.707	BDL	BDL	Not identified	
JBS TP5/1.5m Fill - silty clay, with strong coal tar odours	1.5	<0.001	9.2	0.18	25.1	0.191	BDL	BDL	Not identified	
JBS TP5/2.0m Fill - silty clay, with strong coal tar odours							BDL	BDL	Not identified	
Assessment Criteria										
CH2M Hill RAP ² (maximum of depth dependent values)	5	-	11.8		100	-	42 500	-		
JBS 2010 Site Specific Leachability ³	-	1.7	-	1.162	-	-	-	6.4		

Notes:

1 Bulk sample of same material collected for benchscale immobilisation trial

2 Remediation Acceptance Criteria for Total Concentrations in soil, site specific based on exposure to vapours

3 Site Specific leachability criteria for materials to remain on site

BDL = below laboratory detection limit





Table A9.4: VOC Concentrations in Untreated Soil Samples (units as specified)													
Sample ID	Depth (m)	Total Benzene (mg/kg)	Leachable Benzene (µg/L)	Total Toluene (mg/kg)	Leachable Toluene (µg/L)	Total Ethyl Benzene (mg/kg)	Leachable Ethyl benzene (µg/L)	Total Xylene (mg/kg)	Leachable Xylene (µg/L)	Total 1,3,5 trimethylbenzene (mg/kg)	Leachable 1,3,5 trimethylbenzene (µg/L)	Total 1,2,4 trimethylbenzene (mg/kg)	Leachable 1,2,4 trimethylbenzene (µg/L)
JBS TP1	0.3-0.4	<0.5	<1	<0.5	<1	<1.0	<1	<3.0	<1	<1	<1	<1	<1
JBS TP2	0.4-0.5	<0.5	<1	<0.5	<1	<1.0	<1	<3.0	<1	<1	1.5	<1	2.7
JBS TP2	1.4-1.5	1.4	<1	2.4	<1	26	<1	66	<1	25	340	15	820
JBS TP3	1.7	0.9	<1	0.72	<1	22	<1	31	<1	54	260	36	640
JBS TP3	4-4.2	<0.5	<1	<0.5	<1	<1.0	<1	<3.0	<1	<1	<1	2.2	<1
JBS TP4	0.5	<0.5	<1	<0.5	<1	<1.0	<1	<3.0	<1	<1	<1	<1	<1
JBS TP4	1	<0.5	<1	<0.5	<1	<1.0	<1	<3.0	<1	<1	<1	<1	<1
JBS TP4	1.6-1.7	<0.5	<1	<0.5	<1	<1.0	2.1	<3.0	<1	<1	<1	<1	2.8
JBS TP5	0.5	<0.5	<1	<0.5	<1	<1.0	2.0	<3	<1	<1	19	<1	5.2
JBS TP5	1.5	<0.5	<1	<0.5	35	<1.0	17	8.6	<1	4.1	170	8.5	380
JBS TP5	2.0	<0.5	3.7	<0.5	4.8	<1.0	12	<3	<1	<1	11	<1	31
Assessment Criteria													
CH2M F (maxin depth de valu	num of pendent	1	-	7.9	-	34.8	-	14	-	-	-	-	-
JBS Site Leach		-	8000	-	2880	-	80	-	10 000	-	-	-	-





Table A9.5:		Heavy	Metal Conce	entrations ir	Untreated	Soil Sample	s (units as s	specified)							
Sample ID	Depth (m)	Total Arsenic (mg/kg)	Leachable Arsenic (µg/L)	Total Cadmium (mg/kg)	Leachable Cadmium (µg/L)	Total Chromium (mg/kg)	Leachable Chromium (µg/L)	Total Copper (mg/kg)	Leachable Copper (µg/L)	Total Lead (mg/kg)	Leachable Lead (µg/L)	Total Nickel (mg/kg)	Leachable Nickel (µg/L)	Total Zinc (mg/kg)	Leachable Zinc (µg/L)
JBS TP1	0.3-0.4	30	<0.05	1.1	<0.01	26	<0.01	230	0.1	220	<0.03	20	0.02	260	1.6
JBS TP2	0.4-0.5	13	<0.05	<0.5	<0.01	19	<0.01	80	0.04	220	0.09	26	<0.02	220	1.7
JBS TP2	1.4-1.5	<4	<0.05	<0.5	<0.01	14	<0.01	4	0.04	58	< 0.03	5	<0.02	200	0.5
JBS TP3	1.7	<4	<0.05	<0.5	<0.01	12	<0.01	1	0.02	14	<0.03	2	<0.02	3	1
JBS TP3	4-4.2	5	<0.05	<0.5	<0.01	22	<0.01	6	0.02	24	<0.03	3	<0.02	9	0.2
JBS TP4	0.5	8	< 0.05	<0.5	<0.01	22	<0.01	46	0.1	260	0.05	10	<0.02	4	1.4
JBS TP4	1	9	<0.05	<0.5	<0.01	35	<0.01	51	0.06	61	0.04	3	<0.02	14	0.2
JBS TP4	1.6-1.7	6	<0.05	<0.5	<0.01	7	<0.01	65	0.4	100	0.06	4	<0.02	47	1.4
JBS TP5	0.5	6	<0.05	<0.5	<0.01	18	<0.01	18	0.02	58	0.05	5	<0.02	35	0.3
JBS TP5	1.5	5	<0.05	<0.5	<0.01	23	<0.01	16	0.02	50	<0.03	13	<0.02	93	1.3
JBS TP5	2.0	6	< 0.05	<0.5	<0.01	2.5	<0.01	9	0.02	36	0.03	7	<0.02	27	0.09
Assessment Criteria															
CH2M Hill (maximum c dependent	of depth	1	-	7.9	-	34.8	-	14	-	-	-	-	-		
JBS Site S Leachab		-	8000	-	2880	-	80	-	10 000	-	-	-	-		





ble A9.6 Mean Results of Soil Treatment Trial (units as specified) – Assessment of Compliance to IA 2005/14										
BS/Enviropacific)	Cement ratio	Total B(a)P (mg/kg)	Leachable B(a)P (mg/L) ¹	Total PAHs (mg/kg)	Leachable PAHs (mg/L) ¹	Total TPH C ₁₀ -C ₃₆ (mg/kg)	Leachable TPH C ₁₀ -C ₃₆ (µg/L) ¹	Mean UCS 7-day curing (MPa)	Compliant with IA 2005/14	Waste Classification under IA 2005/14
ol Sample	untreated	1.7	<0.001	51	0.083	500	680	ND	-	-
Material 1 Post 5%	1:20	4.7	<0.001	44	<0.001	360	400	2.18	Y	General Solid
Material 1 Post 12.5%	1:8	3.8	<0.001	34	<0.001	320	280	3.10	Y	General Solid
Material 1 Post 20%	1: 5	3.9	<0.001	43	<0.001	300	290	5.85	Y	General Solid
Material 2 Post 5%	1:20	2.0	<0.001	171	2.6	810	6700	0.35	N	-
Material 2 Post 12.5%	1:8	1.6	<0.001	117	1.9	450	5700	1.00	Y	General Solid
Material 2 Post 20%	1: 5	1.6	<0.001	101	1.5	420	4900	1.55	Y	General Solid
Material 3 Post 5%	1:20	0.8	<0.001	17	0.26	<250	1420	0.13	N	-
Material 3 Post 12.5%	1:8	0.8	<0.001	20	0.33	<280	1300	0.43	Ν	-
Material 3 Post 20%	1: 5	1.7	<0.001	48	0.48	460	1950	0.60	N	-
				Assess	ment Criteri	a				
14, NSW EPA ²	2:1	500	-	13 000	-	-	-	1	-	-
General Solid Waste (DECC 2009)		10	40	200	-	10 000	-	-	-	-
Waste (DECC 2009)	-	23	160	800	-	40 000	-	-	-	-
chability (JBS 2010)	-	-	1.6							
	BS/Enviropacific) ol Sample Material 1 Post 5% Material 1 Post 20% Material 2 Post 20% Material 2 Post 20% Material 3 Post 20% Material 3 Post 5% Material 3 Post 20%	BS/Enviropacific) Cement ratio ol Sample untreated Material 1 Post 5% 1:20 Material 1 Post 5% 1:20 Material 1 Post 20% 1: 5 Material 2 Post 5% 1:20 Material 2 Post 5% 1:20 Material 2 Post 20% 1: 5 Material 3 Post 20% 1: 5 Material 3 Post 5% 1:20 Material 3 Post 5% 1:20 Material 3 Post 20% 1: 5 Material 3 Post 20% 1: 5 Material 3 Post 20% 1: 5	BS/Enviropacific) Cement ratio Total B(a)P (mg/kg) ol Sample untreated 1.7 Material 1 Post 5% 1:20 4.7 Material 1 Post 5% 1:20 4.7 Material 1 Post 20% 1:5 3.9 Material 2 Post 5% 1:20 2.0 Material 2 Post 5% 1:20 2.0 Material 2 Post 5% 1:20 0.0 Material 3 Post 5% 1:5 1.6 Material 3 Post 5% 1:20 0.8 Material 3 Post 5% 1:20 0.8 Material 3 Post 20% 1:5 1.7 Material 3 Post 20% 1:5 1.7	BS/Enviropacific) Cement ratio Total B(a)P (mg/kg) Leachable B(a)P (mg/L)1 ol Sample untreated 1.7 <0.001	BS/Enviropacific) Cement ratio Total B(a)P (mg/kg) Leachable B(a)P (mg/L) ¹ Total PAHs (mg/kg) ol Sample untreated 1.7 <0.001	BS/Enviropacific) Cement ratio Total B(a)P (mg/kg) Leachable B(a)P (mg/L)1 Total PAHs (mg/kg) Leachable PAHs (mg/L)1 ol Sample untreated 1.7 <0.001	BS/Enviropacific) Cement ratio Total B(a)P (mg/kg) Leachable B(a)P (mg/L)1 Total PAHs (mg/kg) Leachable PAHs (mg/L)1 Total Total TPAHs (mg/kg) ol Sample untreated 1.7 <0.001	BS/Enviropacific) Cement ratio Total B(a) (mg/kg) Leachable B(a)P (mg/L) ¹ Total PAHs (mg/kg) Leachable PAHs (mg/kg) Total TPH C ₁₀ -C ₃₆ (mg/kg) Leachable TPH C ₁₀ -C ₃₆ (mg/kg) ol Sample untreated 1.7 <0.001	BS/Enviropacific) Cement ratio Total B(a)P (mg/kg) Leachable B(a)P (mg/L) ¹ Total PAHs (mg/kg) Leachable PAHs (mg/L) ¹ Total TPH C ₁₀ -C ₃₆ Leachable (mg/kg) Mean UCS 7-day (mg/L) ¹ ol Sample untreated 1.7 <0.001	BS/Enviropacific) Cement ratio Total B(a)P (mg/kg) Leachable B(a)P (mg/L)1 Total PAHS (mg/kg) Leachable PAHS (mg/kg) Total TPH Cto-C2s (mg/kg) Leachable (mg/kg) Mean UCS T-day (mg/kg) Compliant with IA 2005/14 ol Sample untreated 1.7 <0.001

Notes 1. Leachability testing by ASLP, unless otherwise specified

2. NSW EPA 'General Immobilisation of Contaminants in Waste – Coal tar Contaminated Waste From Former Gasworks Sites' approval number 2005/14 (IA 2005/14).





Table A9.	7: Mean	Results of Soi	l Treatment Tria	l (units as spe	cified) – As	sessment of	Additional Co	ompound Re	sults			
Sample II) (JBS/Enviropacific)	Cement ratio	Total Naphthalene (mg/kg)	Leachable naphthalene (mg/L) ¹	Total Toluene (mg/kg)	Leachable Toluene (µg/L) ¹	Total Ethyl Benzene (mg/kg)	Leachable Ethyl benzene (µg/L) ¹	Total Benzene (mg/kg)	Leachable Benzene (µg/L) ¹	Total Xylene (mg/kg)	Leachable Xylene (µg/L) ¹
Control Sa	mple	untreated	18	0.059	<0.5	0.083	<0.5	1.5	<0.5	<1	<3	<3
	Material 1 Post 5%	1:20	0.6	0.001	<0.5	<0.001	<0.5	<1	<0.5	<1	<3	<3
JBS TP1/ 0.3-0.4m	Material 1 Post 12.5%	1:8	0.5	<0.001	<0.5	<0.001	<0.5	<1	<0.5	<1	<3	<3
0.3-0.411	Material 1 Post 20%	1: 5	0.4	<0.001	<0.5	<0.001	<0.5	<1	<0.5	<1	<3	<3
JBS TP3/1.7m	Material 2 Post 5%	1:20	120	2.4	<0.5	<1	2.8	70	<0.5	<1	4.7	121
	Material 2 Post 12.5%	1:8	74	1.7	<0.5	<1	1.4	44	<0.5	<1	1.2	80
	Material 2 Post 20%	1: 5	60	1.3	<0.5	<1	<1	30	<0.5	<1	<3	52
JBS	Material 3 Post 5%	1:20	2.7	0.2	<0.5	0.26	<0.5	4.4	<0.5	<1	<3	2.3
TP3/4.0- 4.2m	Material 3 Post 12.5%	1:8	5.0	0.27	<0.5	0.33	<0.5	<1	<0.5	<1	<3	<3
	Material 3 Post 20%	1: 5	16	0.42	<0.5	0.48	<0.5	2.6	<0.5	<1	<3	<3
Assessme	nt Criteria											
IA 2005/	14, NSW EPA ²	2:1	-	-	-	-	-	-	-	-	-	-
General Solid Waste (DECC 2009) -		-	-	-	-	-	-	-	-	-	-	-
Restricted Solid Waste (DECC 2009) -		-	-	-	-	-	-	-	-	-	-	-
Site Specific Leachability (JBS 2010) -		-	-	1.16	-	2880	-	80	-	8000	-	10 000

Notes 1. Leachability testing by ASLP, unless otherwise specified 2. NSW EPA 'General Immobilisation of Contaminants in Waste – Coal tar Contaminated Waste From Former Gasworks Sites' approval number 2005/14 (IA 2005/14).



The results in **Table A9.6** indicate that two of the three materials tested are suitable for treatment by cement stabilisation with the addition of at least 12.5% cement. Material 3, collected from JBS TP3/ 4.0 - 4.3 m, failed to meet the required UCS value, which is likely to be related to the clay content of this material. The results for this material however show that the UCS value increased as the percentage of cement added increased, which suggests that cement stabilisation may be possible using a higher percentage of cement.

Results for the remaining two samples after treatment show full compliance with the requirements of IA 2005/14, and under this order would be suitable for off site disposal to landfill as General Solid Waste at a landfill licenced to receive immobilised material.

With regards to on site reuse, the treated material must be compliant with all the requirements of IA/2005 (to demonstrate contaminants have been stabilised) and the leachable concentrations must be less the site specific leachability criteria. The results in Table A9.6 indicate that only Material 1 complies with all these requirements. As discussed previously Material 3, collected from JBS TP3/ 4.0 - 4.3 m, failed to meet the required UCS value, however this may be rectified using a higher portion of cement. Leachable concentrations of naphthalene in the Material 3 treated samples exceeded site specific leachability criteria, and would therefore be considered unsuitable for reuse on the site. We note that the methods currently accepted for determining leachable concentrations will greatly overestimate leachable concentrations in stabilised material. The Enviropacific report (Appendix I) for the stabilisation trial notes that '...current leachate methods (TCLP, ASLP and MEP are the only leachate tests currently used by regulatory authorities in Australia) have important limitations. For example, each of these methods require particle size reduction to either 9.5 mm (TCLP and MEP) or 2.4 mm (ASLP), which effectively contravenes the assessment of encapsulated wastes as the integrity of the monolithic structure is compromised, and each of these methods employ vigorous end-over-end agitation of the sample. In fact in AS4439.3- 1997 (ASLP) the Scope states that "The procedure is not applicable to encapsulated wastes which cannot be reduced to the specified maximum particle size without breaking the integrity of encapsulation". For this reason, in the General IA, the DECCW is in effect relying on the UCS measurement as an indicator of the stability of cement stabilised waste, whilst still requiring B(a)P leachability (TCLP) for cement stabilised samples to be below the Waste Classification Guidelines criteria (and from previous EPS experience with other gasworks projects B(a)P leachability has typically been non-detectable in both the untreated and cement stabilised samples using TCLP).

If a more appropriate leach test (e.g. a diffusion-based or column leach test on a moulded/monolithic sample) was adopted for assessing the leachability of cement stabilised materials destined for on-site placement, the optimum UCS required to minimise contaminant leachability could be more accurately determined, and a different leachability data set may result, that might be more appropriately applied to acceptance criteria for on-site placement of cement stabilised material.'

The results of the treatment trial suggest that some cement stabilised soils may be suitable for on-site reuse. The limited sample size suggests that material containing total naphthalene concentrations over 50 mg/kg may not be suitable for on site reuse following stabilisation, or use of column leach tests in addition to ASLP testing may be required to determine the leachable concentrations in cement stabilised material. A column leach test, although not an approved method, would avoid grinding of the treated material and be more representative of leaching conditions that may occur on site. TCLP testing would still likely be required for assessment of the material against IA 2005/14, and determining the waste classification of the treated material if off site disposal is required.



At the request of the Site Auditor samples of untreated material were also analysed after the materials had been sorted (*i.e.* oversize particles removed) and prior to the addition of water and/or cement. Full tabulated pre and post treatment analytical results are presented in **Appendix J.**

As expected, for each of the bulk samples supplied, the range of total concentrations in the subsampled material shows minor variation. Generally the total concentrations in the three subsample prepared from each bulk material were within 20 percent of each other.

With the exception of B(a)P, total concentrations generally showed only minor reductions between the pre and post treatment samples. The slight reduction is considered to be the result of the addition of cement to the material, essentially diluting the original concentrations. Contaminant reductions above those noted were not expected as the trialled treatment method is not intended to remove contamination only reduce its mobility.

To assess the ability of the treatment method to reduce mobility the leachable concentrations were compared in the pre and post treatment samples. It is noted that pre treatment leachable concentrations were determined using the TCLP method to assess the likely waste classification of these materials assuming no treatment occurs. Leachable concentrations in the post treatment samples were determined using the ASLP method, to assess the suitability of cement stabilised material to be reinstated on site. The following paragraphs generally discuss the variation in leachable concentrations between pre and post treatment samples as an indication of the likely performance of the material, although readers should be aware of the different leachability methods used in determining these results.

BTEX compounds and naphthalene were noted as the main contaminants of concern in groundwater migrating off site. The following was noted in relation to leachable concentrations in the samples untreated and treated from the trial.

In general the leachable concentrations of all BTEX compounds, where detected, were reduced by an order of magnitude in the treated material samples. It is however noted that all leachable BTEX concentrations in the samples of untreated material were less than the site specific leachability criteria.

For naphthalene mixed results were obtained, the leachable concentrations in Material 1 (JBS TP1/0.3-0.4m) were less than the laboratory detection limit in both the untreated and treated material. For Material 2 (JBS TP1.7 m), naphthalene concentrations on average reduced by only 1 mg/L between the pre and post treatment samples. This may be due to a number of factors including the elevated total concentrations of naphthalene, the use of the ASLP method (rather than column leach tests) as discussed above. In Material 3 (JBS TP3/4.0-4.2 m) the leachable concentrations in the treated material were an order of magnitude less than concentrations detected in the untreated material. It is noted that leachable naphthalene concentrations in all Material 3 samples, both untreated and treated exceeded the site specific leachability criteria. These results suggest that where material is being treated by cement stabilisation for on site reuse additional refinement of the treatment process is required to reduce the ASLP determined leachable concentration. Alternately consideration should be given to use of method to determine the leachable concentration.

While B(a)P was not listed as a contaminant of concern in groundwater it is noted that the leachable concentrations in all untreated and treated samples were less than the laboratory detection limit, consistent with the results discussed in **Section A9.3**.



A9.5 Water Treatment Trial

The methods and results of the water treatment trial conducted on the site were documented in the JBS Letter Report '*Groundwater Treatment Trial, Former Macdonaldtown Gasworks*', dated 05 August 2010 Reference 40913 - 15534. A copy of that letter is included as **Appendix C**.

The results of the water treatment trial are summarised in **Tables A9.8** and **A9.9** below. Overall it is noted that samples of the treatment system effluent (*i.e.* all 'Sample 4' results, collected post GAC filter) were reported to contain very low concentrations of all contaminants of concern at the site. Based on these results it appears that a water treatment plant could be used as part of the remediation process to enable discharge of collected groundwater as follows:

- To stormwater following receipt of relevant approvals from City of Sydney Council; or
- To sewer should a 'trade waste agreement' be entered into with Sydney Water.

Additionally the results indicate that the treated effluent was of a suitable quality for reinjection into the subsurface. The feasibility of this option, is however, uncertain, given the required discharge rates during remediation are likely to exceed the infiltration potential of the clay soils underlying the site. This option would also require licensing by the NSW Office of Water.





Table A9.8:	Summary of Water Tre	eatment Tria	il Results – V	OCs, TPH and	Heavy Metals									
							lyte							
			VOCs	r		РН		Heavy Metals						
Sample ID	рН	Chloro- form	Bromo- dichloro- methane	Dibromo- chloro- methane	C ₆ -C ₉	C ₁₀ -C ₃₆	As	Cd	Cr	Cu	Pb	Hg	Ni	Zn
MW04S	Pumped groundwater holding tank	21	25	18	<10	6,200	<1	<0.1	<1	<1	<1	<0.5	1	110
Sample 1 – Run 1	Influent (15 mins)	14	8.9	2.7	<10	1,100	<1	<0.1	<1	<1	<1	<0.5	1	110
Sample 2 – Run 1	Post oil-water separator / pre sand filter (15 mins)	9.8	5.8	1.7	<10	170	<1	<0.1	<1	1	<1	<0.5	2	100
Sample 3 – Run 1	Post sand filter/ pre GAC (15 mins)	12	7.6	2.2	<10	3,210	<1	<0.1	<1	10	<1	<0.5	8	63
Sample 4 – Run 1	Post GAC Effluent (15 mins)	<1	<1	<1	<10	<250	16	<0.1	<1	<1	<1	<0.5	<1	1
Sample 1 – Run 2	Influent (30 mins)	12	2.1	2.1	<10	3,600	<1	<0.1	<1	3	<1	<0.5	7	170
Sample 2 – Run 2	Post oil-water separator / pre sand filter (30 mins)	11	6.6	2.0	<10	322	<1	<0.1	<1	4	<1	<0.5	9	91
Sample 3 – Run 2	Post sand filter/ pre GAC (30 mins)	9.7	6.3	2.0	<10	190	<1	<0.1	<1	7	<1	<0.5	8	48
Sample 4 – Run 2	Post GAC Effluent (30 mins)	<1	<1	<1	<10	<250	17	<0.1	<1	<1	<1	<0.5	<1	<1
Sample 1 – Run 3	Influent (40 mins)	9.6	5.9	1.8	<10	5,700	<1	<0.1	<1	<1	<1	<0.5	2	160
Sample 2 – Run 3	Post oil-water separator / pre sand filter (40 mins)	10	5.5	1.7	<10	<250	<1	<0.1	<1	<1	<1	<0.5	2	140
Sample 3 – Run 3	Post sand filter/ pre GAC (40 mins)	11	5.2	1.6	<10	1230	<1	<0.1	<1	4	1	<0.5	2	49
Sample 4 – Run 3	Post GAC Effluent (40 mins)	<1	<1	<1	<10	<250	24	<0.1	<1	<1	<1	<0.5	<1	<1





able A9.9:	Summary of Wa	ter Treat	ment Tria	l Results	- PAHs			Δn	alyte - PA	\He						
Sample ID	Component or sampling stage (time after system commencement)	Naphthalene	Acenaphthylene	Acenaphthene	Fluorene	Phenanthrene	Anthracene	Fluoranthene	Pyrene	Benzo(a)anthracene	Chrysene	Benzo(b+k)fluoranthene	Benzo(a)pyrene	Indeno(1,2,3-c,d)pyrene	Dibenzo(a,h)anthracene	Benzo(g,h,i)perylene
MW04S	Pumped groundwater holding tank	5.6	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1
Sample 1 – Run 1	Influent (15 mins)	2.2	<0.1	0.7	0.5	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1
Sample 2 – Run 1	Post oil-water separator / pre sand filter (15 mins)	0.3	<0.1	0.4	0.4	0.4	<0.1	0.2	0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1
Sample 3 – Run 1	Post sand filter/ pre GAC (15 mins)	0.2	<0.1	0.3	0.2	0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1
Sample 4 – Run 1	Post GAC (15 mins)	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.02	<0.01	<0.01	<0.01	<0.01
Sample 1 – Run 2	Influent (30 mins)	3.4	<0.1	0.5	0.4	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1
Sample 2 – Run 2	Post oil-water separator / pre sand filter (30 mins)	0.2	<0.1	0.5	0.5	1	0.2	0.7	0.4	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1
Sample 3 – Run 2	Post sand filter/ pre GAC (30 mins)	0.2	<0.1	0.4	0.4	0.5	<0.1	<0.1	<0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1
Sample 4 – Run 2	Post GAC (30 mins)	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.02	<0.01	<0.01	<0.01	<0.01
Sample 1 – Run 3	Influent (45 mins)	3.8	<0.1	0.8	0.6	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1
Sample 2 – Run 3	Post oil-water separator / pre sand filter (40 mins)	0.2	<0.1	0.6	0.5	0.1	<0.1	0.1	<0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1
Sample 3 – Run 3	Post sand filter/ pre GAC (40 mins)	0.2	<0.1	0.6	0.5	0.6	0.1	0.1	0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1
Sample 4 – Run 3	Post GAC (40 mins)	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.02	<0.01	<0.01	<0.01	<0.01



A9.6 Pump Tests

The groundwater wells pumped to generate water for the treatment trials were also subject to small scale pump tests, conducted in line with the method provided in MacDonald A, Barker J and Davies J (September 2008) '*The bailer test: a simple effective pumping test for assessing borehole success*' in <u>Hydrogeology Journal.</u>

The following locations were subject to pump tests:

- MW04S;
- MW07S;
- MW37S; and
- MW42S.

Data loggers were used to record the drawdown in these wells during the pumping and recovery phases of the tests. Line graphs of the raw data collected from each well over the duration of the pump test are presented in **Figures A2** to **A3**, as time versus height of water above the logger.

Based on this inspection of the raw data, no further analysis was conducted on the results from MW04S and MW07S. The plots from these two locations show that both wells were unable to sustain pumping rates of 1 - 2.4 Litres per minute (L/min), with both running dry over the duration of pumping. While these results reflect the spatial variability in the volume of perched groundwater present under the site, and permeability of the formation, they were considered to not provide meaningful data for remediation planning.

The results from MW37S and MW42S were used to estimate the hydrologic properties of fill material underlying the site. The pump test data from both locations were used to estimate the transmissivity of the screened formation using three methods of analysis as follows:

- Papodopulos and Copper (1967) –using the active pumping data only;
- Cooper and Jacob (1946) using the active pumping data only; and
- Theis Recovery Method (1946) using the well recovery data only.

The methods selected have been assessed as suitable for analysis of constant discharge, single well tests (*i.e.* no use of observation wells during the pump test) in unconfined aquifers (Kruseman & De Ridder, 2000). Kruseman & De Ridder (2000) also indicates that as storage in the well may influence drawdown at the commencement of pumping and recovery phases of the test, then only the latter stages of data should be used for curve matching.

Curve matching and data analysis were undertaken using the AquiferWin32 software package. The transmissivity values calculated using all three tests are summarised in **Table A9.10** below, with the corresponding AquiferWin32 output files and curve matching analysis provided in **Appendix I**.

The hydraulic conductivity of the formation surrounding the pumped wells was estimated from the calculated transmissivity values using the following equation:

- K = T/b Where T = aquifer transmissivity
 - b = aquifer thickness (assumed to be the length of the saturated screen interval in the well);
 - K = hydraulic conductivity of the formation.



	SWL*	Screen	Trans	mittivity (m²	.s⁻¹)	Mean transmissivity	Hydraulic conductivity	
Well ID	(m below TOCª)	Interval (m below TOC)	Papodopulos & Copper (1967)	Jacob	Theis Recovery Method (1946)	(m ² .s ⁻¹)	(m.s ⁻¹)	
MW37S	0.931	1.5-4.5	9.89 x 10 ⁻⁶	1.23x 10 ⁻⁶	3.47 x 10 ⁻⁵	1.55 x 10 ⁻⁵	5.09 x 10 ⁻⁶	
MW42S	1.19	1.5-4.5	9.70 x 10 ⁻⁶	1.32 x 10 ⁻⁶	4.88 x 10 ⁻⁵	1.99 x 10 ⁻⁵	6.65 x 10 ⁻⁶	
Notes:	* (Standing Wa	ter Level					

com the values estimated in Table A9 10, hydraulic conductivi

From the values estimated in **Table A9.10**, hydraulic conductivity values for the screened intervals within the wells tested range between $5.09 \times 10^{-6} \text{ m.s}^{-1}$ and $6.65 \times 10^{-6} \text{ m.s}^{-1}$.

Very few published values are available for fill layers, given the inherent variability of groundwater flow characteristics in these materials, it is, however, noted that the estimated values of hydraulic conductivity are in the range of values listed in Freeze and Cherry (1979), for silts, sand and fine grained sand.

The hydraulic conductivities reported in **Table A9.10** are equivalent to an order of magnitude less than the average value of 1.4×10^{-5} m.s⁻¹ reported for clay soils in (SKM 2006). In relation to the slug test determined conductivity value, it is noted that the SKM report noted that '...based on the measured extent of the shallow and deep groundwater plumes migrating from the site, the permeability rates estimated from the slug tests appear to be an order of magnitude greater than actual rates'. The JBS calculated conductivity values therefore, appear to be consistent with rate of off-site migration of contaminated groundwater.

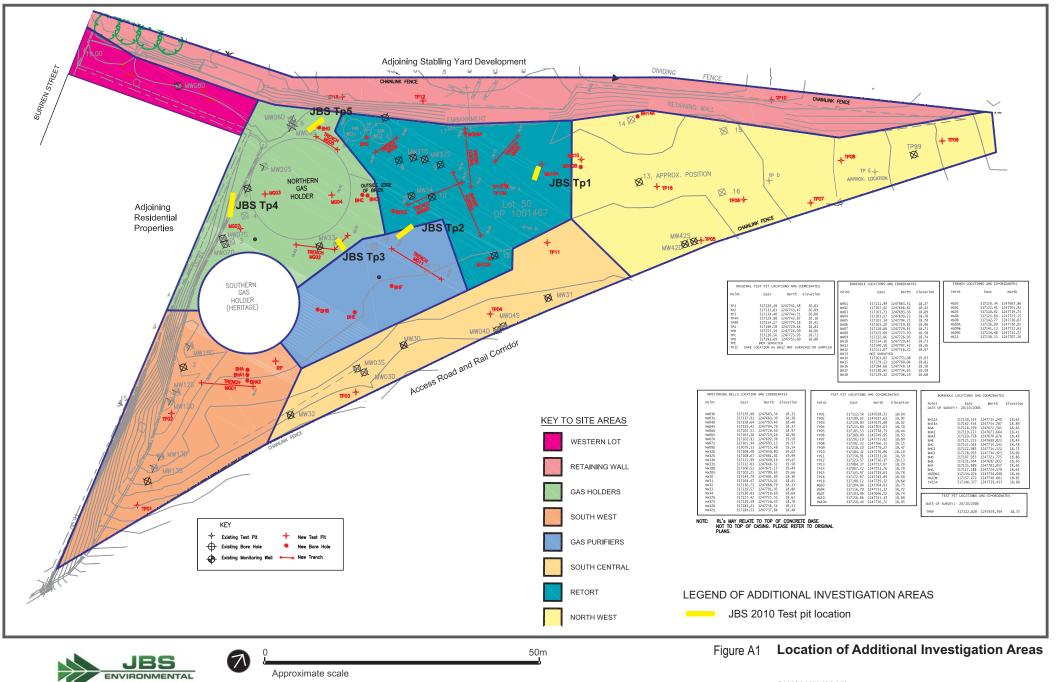
Adopting a maximum hydraulic gradient of 3.1×10^{-2} the SKM report estimated flow velocities in shallow groundwater to be on the order of 6.2 and 13.7 metres per year. Using the same hydraulic gradient value, the conductivities reported in **Table A9.10** equate to flow velocities between 5 and 6.5 metres per year.

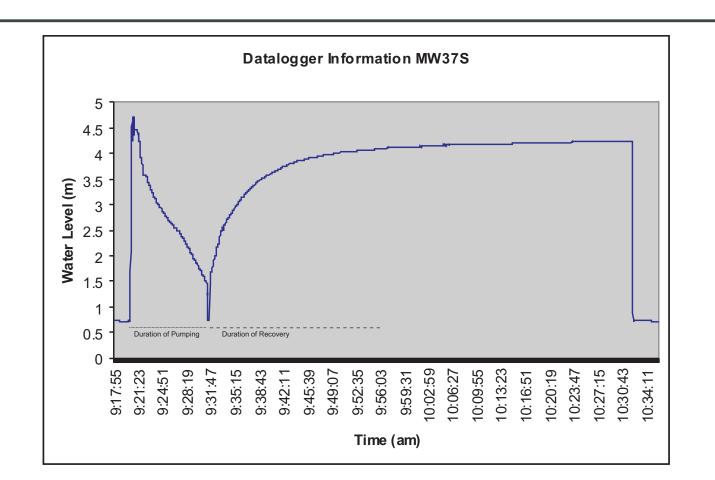
The difference in estimated hydraulic conductivities may be attributable to the use of a pump test in the recent work by JBS, compared to slug tests by SKM in 2006. The pump test undertaken by JBS enabled estimates of conductivity to be made from both the pumping and recovery periods, while the slug test only allows for assessment of recovery.

Given that two of the four shallow wells ran dry over the duration of pumping, it is considered that the values provided in **Table A9.10** are likely to represent the upper end of hydraulic conductivities in fill on the site.



Figures A1 to A3





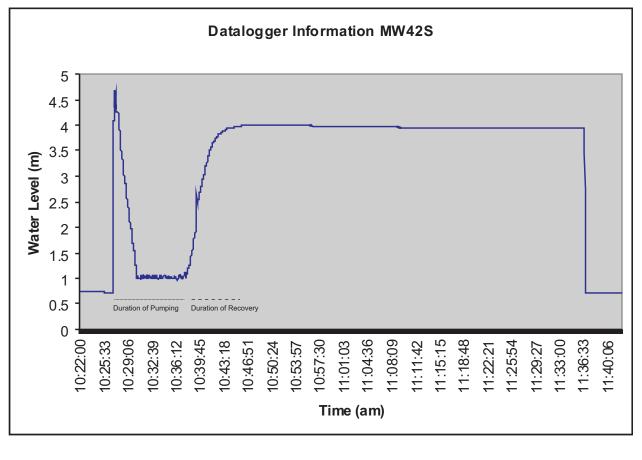




Figure A3 Raw Pump Test Data MW37S and MW42S

Former Macdonaldtown Gasworks Burren Street, Erskineville Job Reference: 40913 R08

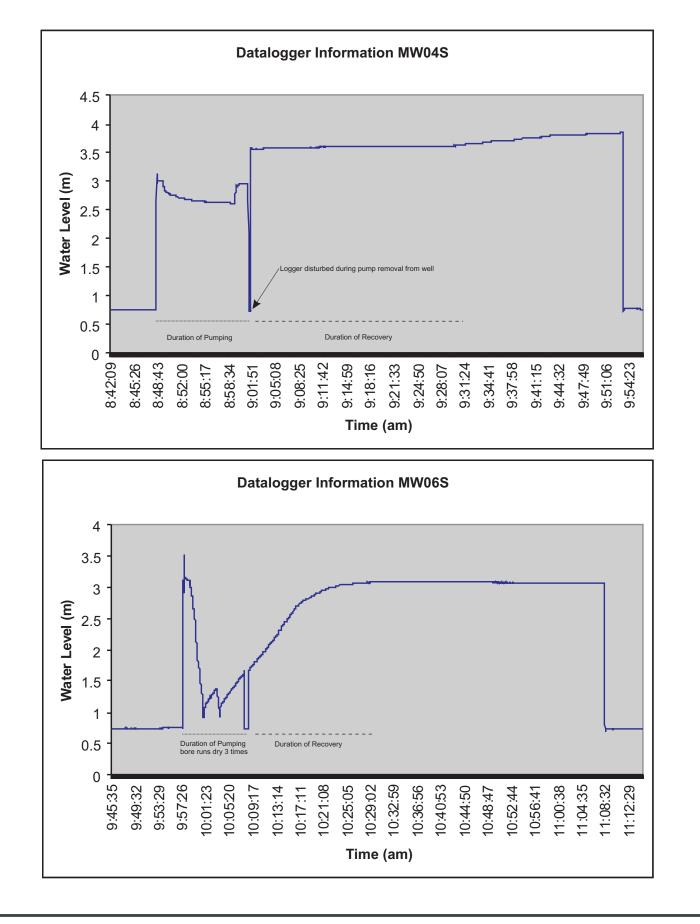




Figure A2 Raw Pump Test Data MW04S and MW06S

Former Macdonaldtown Gasworks Burren Street, Erskineville Job Reference: 40913 R08



Appendix B

`Site Specific Leachability Criteria, former Macdonaldtown Gasworks, Burren Street Erskineville' JBS Environmental Pty Ltd (2010a)





JBS40913-15507

23 August 2010

John Dawson Project Director Incoll Management Pty Ltd Level 1, 73 Miller Street North Sydney NSW 2060 (Sent Via email: jdawson@incoll.com.au)

Development of Site Specific Soil Leachability Criteria – Former Macdonaldtown Gasworks, Burren Street, Erskineville, NSW

Dear John,

JBS Environmental Pty Ltd (JBS) was commissioned by Incoll Management Pty Ltd (Incoll), to undertake additional environmental assessment works to inform the revision of the remedial strategy prepared for the former Macdonaldtown Gasworks site, on behalf of the site owner, Rail Corporation NSW Environment Projects Unit (RailCorp). This letter has been prepared as part of those works and documents the derivation of site specific criteria for leachable concentrations of contaminants in soil. These are proposed to be used for the proposed remediation of the site.

The site is located between Erskineville and Macdonaldtown railway stations. The site is roughly triangular in shape, being part of the area commonly referred to as the Macdonaldtown Triangle. The site location is shown in **Figure 1**.

A Remedial Action Plan (RAP) has been prepared for the site and is documented in CH2M Hill (December 2007) '*Remedial Action Plan, Former Macdonaldtown Gasworks – Burren Street, Erskineville, NSW*. Based on review of the RAP it is considered that while a range of technologies may be applicable to the site, the remedial program is likely to be a combination of the following methods:

- Dewatering of impacted areas to enable excavation of fill / soil as required;
- On-site treatment of water generated during dewatering for groundwater recharge, discharge to stormwater or discharge to sewer, as appropriate;
- Removal / excavation of free tar as required for disposal off site to an appropriately licenced landfill or treatment facility;
- Excavation of impacted soil and for treatment by stabilisation, thermal desorption or other appropriate method; and
- Reuse of stabilised material on the Macdonaldtown site, if suitable, otherwise disposal off site to an appropriately licenced landfill.

In order to implement such a remedial approach, in addition to health based assessment criteria for protection of future site users, it will also be necessary to derive site specific acceptance criteria for leachable concentrations in soil that are considered suitable for protection of the environment. Once derived it is considered these site specific criteria can then be applied (in combination with human health based soil criteria as total concentrations) at two levels;

- Initially on *in-situ* impacted materials to assess whether excavation / remediation is required for protection of environmental values; and
- Secondly on treated materials to assess whether they are suitable for reuse on the site.

This letter has been prepared to document the derivation of the site specific criteria for leachable concentrations in treated material for use during remediation of the site.

1. Site Background

Calculation of site specific leachability criteria requires the input of site specific data relating to the hydrogeological conditions and contamination present. This Section provides a summary of this information and is based on a review of the following documents prepared for the site:

- Rail Services Australia (November 1999) 'Eveleigh Gasworks Site History';
- CH2M Hill (December 2001) 'Soil and Groundwater Investigations of the former Gasworks Area and Offsite';
- Sinclair Knight Merz (April 2006) 'Macdonaldtown Triangle (Former Gasworks Site) Human Health and Ecological Risk Assessment';
- CH2M Hill (March 2007) 'Delineation & Characterisation Sampling and review of Remedial Options'; and
- CH2M Hill (December 2007) 'Remedial Action Plan'.

A summary of the subsurface profile encountered on the site is provided in **Table 1**.

2. Hydrogeology

The review of the hydrogeological conditions at the site included in the RAP (CH2M Hill, 2007) states that the groundwater exists as a shallow perched groundwater 'layer ' and a deep bedrock 'layer'. The shallow groundwater, encountered as shallow as 1m below ground level in some areas, was reported to be present within fill materials and silty clay overlying the mottled clay layer as describes in **Table 1**, while the deeper groundwater was reported to exist within the Ashfield Shale bedrock under semiconfined conditions.

A summary of estimates of the hydrogeological properties of the subsurface is provided in **Table 2**, and is based on data provided in SKM (2006).

All previous reports acknowledge that while shallow groundwater underlying the site appears to be restricted to the fill and clay layers, and deep groundwater to the shale bedrock, it is possible there may be some interconnectivity between the two water bearing zones given the apparent similar direction of flow gradient.

Layer	Observed Depth	Nature of Material		Nature of Impact	
	(m below ground level)		Free Tar ¹	Tarry Soils ²	Dark stained impacts ³
		Ash and coke gravels to 0.5m depth across much of the site	✓ Limited to immediate vicinity of tar wells and gas works pipes	✓ Limited to areas of former gasworks footprints	-
	Reworked clay 0.5m to 1.5m depth across the majority of the site	✓ Limited to immediate vicinity of tar wells and gas works pipes	✓ Limited to areas of former gasworks footprints	-	
Fill	Fill 0 to 3.5	Sands and gravels 0.5m to 1.5m depth in north-east, south central and gas purifier areas	 ✓ Limited to immediate vicinity of tar wells and gas works pipes 	✓ Limited to areas of former gasworks footprints	-
		Gravelly Sand and Clay with minor ash to depths of 3.5m in the south west area of the site	-	✓ Limited to areas of former gasworks footprints	-
		Gravelly sand and demolition waste in the northern retaining wall and inside annulus of northern gas holder	-	✓ Limited to areas of former gasworks footprints	-
Silty clay	1.5-2.5	Saturated silty clay layer present underlying fill across the majority of the site	✓ near tar wells and northern gasholder	✓ near tar wells and northern gasholder	-
Red/grey mottled clay	2.5 to 4.0-6.0	Highly plastic, stiff to very stiff, moist and consistent with red podzolic soil	✓ near tar wells and northern gasholder	\checkmark	✓ under southern gasholder
Shale	4 m onwards	Underlies natural clays and grades from extremely weathered to moderately weathered at 10 m depth. Fractures assessed to be common beyond depths of 6m.	✓ near tar wells and northern gasholder	-	✓ under southern gasholder

 Table 1:
 Summary of Subsurface Profile and Contamination Impacts

Notes: 1. 'Free tar' as defined in RAP (CH2M Hill, 2007) is 'soil impacted by free tar, consisting of soil and fill material impacted to a high degree with black ooze, highly odourous, liquor type material';

2. 'Tarry soils' as defined in RAP (CH2M Hill, 2007) is 'soil and fill materials with minor tar impacts and moderate odours'

3. 'Dark Stained impacts' as defined in RAP (CH2M Hill, 2007) is 'dark brown to black staining in the deep soils and Weathered Shale within the soil pores and shale fracture zones underneath the Southern Gasholder. The material as moderately odourous';

✓ = present in layer

not present in layer

Parameter	Value	Details	Sources				
Flow direction	South to south- easterly	0 0	tion in shallow groundwater flow due to surface obstructions				
	1.4 x 10 ⁻⁵ ms ⁻¹	Shallow well					
Permeability	1.0 x 10 ⁻⁵ to 3.0 x 10 ⁻⁵ ms ⁻¹	Deep wells	Slug test conducted by SKM in 2005				
Hydraulic Gradient	3.1 x 10 ⁻²	Between shallow wells MW17S and MW12S					
	1.3 x 10 ⁻²	Between shallow wells MW20S and MW03S	Groundwater gauging by SKM in 2005				
Flow velocities	6.2 to 13.7 m/yr	Shallow wells					
Averaging Time – Threshold ¹	12.2 to 36.5 m/yr	Deeper wells	Determined by SKM from gradient and permeability data				

Table 2:	Summary of Hydrogeolgical Data Available for the site
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3. Groundwater Quality

The most recent round of groundwater sampling completed at the site was conducted by SKM in 2005 (SKM, 2006). The maximum concentrations of contaminants exceeding the assessment criteria in the 'on site' wells is summarised in **Table 3**. In general the maximum concentrations in samples collected from the shallow wells were less than concentrations in samples collected from the deeper wells.

	Criteria	Shall	ow Groundwat	er	Dee	ep Groundwate	er
Analyte	ANZECC 2000	Concentration Range	Well ID - highest concentration	Site Area	Concentration Range	Well ID - highest concentration	Site Area
Cd	0.2	nd - 2.6	MW13s	Southwest	nd - 1.5	MW06d	Gasholders
Cr(total)	-	nd - 15	MW04s	South Central	nd - 7	MW04d	South Central
Cu	1.4	nd - 220	MW42s		0.001 - 208	MW42d	Northeast
Pb	3.4	nd - 174	MW42s	Northeast	nd - 140	MW03d	South Central
Ni	Ni 11		MW04s	South Central	nd - 92	MW36d	Offsite
Zn	8	0.033 - 1,570	MW13s	Southwest	0.015 - 869	MW42d	Northeast
Cyanide (total)	7	0.02 - 0.479	MW20s	Gasholders	nd - 14.9	MW03d	South Central
Benzene	950	nd - 704	MW07s	Gasholders	nd - 14,000	MW03d	South Central
Toluene	-	nd - 117	MW07s	Gasholders	nd - 792	MW03d	South Central
Ethylbenzene	-	nd - 213	MW07s	Gasholders	nd - 317	MW03d	South Central
Total Xylenes	550 (o & p)	nd - 417	MW07s	Gasholders	nd - 5,010	MW03d	South Central
Total PAHs	16 (naphthalene)	nd - 1,677 (naphthalene 1,460)	MW07s	Gasholders	nd - 4,208 (naphthalene 3,840)	MW07d	Gasholders

Table 3: Summary of Groundwater Contamination

Note: "nd" is 'Non Detect', or less than the laboratory Limit of Reporting (<LOR). All concentrations in $\mu g/L$

Water quality parameters measured in samples from the site in 2005 indicated that groundwater underlying the site was of low salinity, 'slightly variable' pH and highly oxygenated (SKM, 2006) which was considered to be indicative of recharge occurring primarily through rainfall infiltration.

With respect to heavy metals, elevated concentrations of cadmium, copper, nickel and lead and zinc were detected in both the deep and shallow wells located along the northern site boundary. The discussion provided in the SKM report indicated that elevated heavy metals concentrations were representative of background conditions. The only exception noted was a zinc concentration of 1570 μ g/L in the sample from MW13S, which is located in the south-western portion of the site and was the highest zinc concentration detected in the available dataset.

The highest concentrations of Total Petroleum Hydrocarbons (TPH) $C_6 - C_9$, benzene, toluene, ethylbenzene and xylene (BTEX) and volatile organic carbons (VOCs) on site were detected in the deep wells located in the vicinity of the southern gasholder and central southern site boundary at wells MW03D, MW04D, MW07D and MW12D.

Elevated concentrations of TPH $C_{10} - C_{36}$, polycyclic aromatic hydrocarbons (PAHs), primarily being naphthalene, were generally detected in the deep wells located in the vicinity of the southern gasholder and central southern site boundary at wells MW03D, MW04D and MW07D. It is noted that speciated 'indicator' compounds have been identified in each sample where significant levels of TPH have been reported.

Phase separated hydrocarbons were not observed in any wells forming the monitoring well network.

Based on the available data it is considered that the primary levels of contaminants of concern in groundwater at the site are BTEX and naphthalene. Elevated TPH concentrations are consequent of elevated concentrations of BTEX and PAH compounds.

4. Contaminant Hydrogeology

Based on the results of laboratory analysis of groundwater samples collected from on site and off site wells, the SKM (2006) report stated that:

'The shallow plume appears to begin near the northern boundary of the Former Cleaning Shed and Gasworks areas and extend in a south-west direction of some 75m. The data indicate that the downgradient edge of the plume is located at the East Hills Line at the southern edge of the site boundary. The lateral extent of plume appears to be confined in the west to the sewer main located adjacent to the rear boundary of the residential properties, while to [sic] the plume is estimated to extent 50m to the east of the former tank area.

The extent of the middle to heavy-end hydrocarbon plume in the deeper aquifer appears to be larger than the shallow aquifer. While the northern, eastern and western boundaries of the plume are similar to the shallow plume, the down-gradient extent of the plume appears to cover a distance of some 160m from the former tar tank area, with its edge near the southern boundary of railway land along Railway Parade. The data indicate that the deep aquifer plume is located entirely on railway owned [sic] land.'

In relation to the estimated flow velocities for the site, as summarised in **Table 2**, the RAP CH2M Hill, 2007) notes that the flow velocity values provided in SKM 2006, do not correlate with the measured lateral extent of the plume, based on gasworks operations commencing more than 100 years prior. It is also noted that the SKM report states that, based on the measured extent of the shallow and deep groundwater plumes migrating from the site, the permeability rates estimated from the slug tests appear to be an order of magnitude greater than actual rates.

5. Site Specific Environmental Investigations for Leachable Soils Concentrations

In preparing the revised remediation strategy for the site, it is proposed that both the total and leachable concentrations of the contaminants of concern are assessed.

Acceptable total concentrations of contaminants in site soils have been set on human health based criteria in SKM (2006).

The principal potential main environmental exposure pathway for the site is groundwater. Assessment criteria for leachable concentrations of contaminants in soil should be consistent with groundwater ecological criteria. These criteria are considered not appropriate for direct comparison to laboratory measurement of soil leachability. The laboratory testing method for leachable concentrations of soil constituents involves an extended period of tumbling the sample in the media to be analysed. This is highly unlikely to ever be replicated in the environment. Additional correction factors need to be considered to allow comparison of soil leachabilities to groundwater based ecological protection criteria.

Leachability criteria determined for the site should be based on the available criteria used for protection of groundwater resources and incorporate a dilution attenuation factor (DAF). A range of DAF calculation methods currently exist and account for either:

- The dilution of contaminated leachate that occurs as it reaches a water bearing zone which is assumed to be unimpacted, as per methods published by the United States Environmental Protection Agency (USEPA, 1996) and the Center for Research in Water Resources (CRWR, 2003); or
- The dilution of contaminated groundwater that occurs within the overall catchment prior to discharge at the receiving water body. Dilution occurs by mixing with groundwater present across the remainder of the catchment. The basis and calculation of this factor is detailed in the method published by the Commonwealth Scientific and Industrial Research Organisation (CSISO, 2009).

6. Background

The method used to determine the site specific leaching criteria is as follows:

- 1. Review of the site and surrounding area to identify the nearest potentially sensitive ecological receptor as associated environmental values, as described in **Section 6.1**;
- 2. Determine Groundwater Investigation Levels (GILs) for the site by consideration of the environmental values of the surrounding catchment / nearest potentially sensitive receptor, as described in **Section 6.1**;
- 3. Estimate an approximate DAF value based on the likely discharge of groundwater sourced from beneath the site to the nearest potentially impacted environmental receptor, as described in **Section 6.3**;
- 4. Calculate the site specific criteria for leachable concentrations in treated material by multiplying the GIL by the DAF for each of the groundwater contaminants of concern, as described in **Section 6.4**.

6.1. Nearest Potential Sensitive Ecological Receptor

The NSW DEC (2007) '*Guidelines for the Assessment and Management of Groundwater Contamination*' requires groundwater investigation levels to be developed to protect environmental values of the surrounding catchment area. This includes consideration of current and potential future uses of groundwater and related ecosystems.

The site lies within the Alexandra Canal sub-catchment of the Cooks River catchment boundary as established by DECCW. The interim water quality objectives established by DECCW for tributaries within the Cooks River catchment¹ include:

- Aquatic ecosystem;
- Visual amenity;
- Secondary contact recreation;
- Primary contact recreation (for achievement in 10 years or more); and
- Aquatic foods to be cooked before eating (for achievement in 5-10 years).

Groundwater discharges from the site need to be protective of these water quality objectives. These beneficial uses will occur at groundwater discharge point nearest to the site, which is most likely the Munni Street Creek which discharges into Alexandra Canal. Shallow groundwater migrating off site may flow directly into the Munni Street Creek or enter piped stormwater drains that flow into the Creek, located approximately 50 m from the site, along Railway Parade to the south.

Of the water quality objectives listed for the overall Cooks River catchment, only 'protection of aquatic ecosystems' are considered applicable to the nearest surface water discharge point (i.e. Munni Street Creek then Alexandra Canal), noting that:

- Visual amenity has been severely degraded below the 'pristine' condition of this waterway;
- Secondary contact recreation is unlikely to occur given the degraded visual amenity; and
- Primary contact activities and consumption of aquatic foods are also unlikely to occur.

Aquatic ecosystems at the discharge point have been adopted as the environmental values requiring protection from groundwater discharging from the site. The GILs applicable to this value are summarised in **Table 3** and are based on ANZECC/ARMCANZ (2000) water quality trigger values for protection of 95 % of species in marine water.

6.2. Determination of DAF Values

When leachate from soil reaches a water bearing zone, it is mixed with the existing water present in that saturated layer and is diluted as it becomes groundwater. Once part of the groundwater system, it is again diluted at the discharge point, where it becomes mixed with the groundwater discharge from the remainder of the catchment.

In the derivation of soil clean-up criteria for a site that allows for protection of ecological values a DAF is used to account for these processes.

¹ Source: <u>http://www.dec.nsw.gov.au/ieo/CooksRiver/caag.htm</u>

Several methods exist for calculation of the DAF value. A review of the various methods was undertaken as part of this assessment. DAF calculation methods described in the following documents are the most widely used and/or most appropriate methods for use at the site:

- USEPA Soil Screening Level (SSL) Method (USEPA 1996);
- Center for Research in Water Resources, University of Austin Texas (2003); and •
- Method described in CSIRO (2009).

DAF values calculated for the site using each of the listed methods are discussed in the following sections.

USEPA Soil Screening Level (SSL) Method (USEPA 1996)

USEPA Soil Screening Level (USEPA, 1996) provides a default DAF value of 20 for sites occupying an area greater than 0.5 acres, however states that site specific DAF can be used in various options for calculating impact to groundwater soil cleanup criteria, including calculation of the Leachate Criterion (LC). A site specific DAF is calculated using Equation 1 below (taken from the USEPA 1996 guidance document Equation 37). Equation 1 requires a value for the mixing zone depth in the aquifer, which is calculated using Equation 2 (taken from the USEPA 1996 guidance document Equation 45).

$$DAF = 1 + (Kid/IL)$$
 (Equation 1)

Where	i	=	gradient (m/m)
	d	=	mixing zone depth (m), calculated below (Equation 2)
	Ι	=	infiltration rate (m/yr)
	L	=	length of area of concern parallel to ground water flow (m)
	К	=	aquifer hydraulic conductivity (m/yr)

 $d = (0.0112L^2)^{0.5} + d_a \{1 - \exp[(-LI)/(Kid_a)]\}$

(Equation 2)

Where

da

aquifer thickness (m) - (USEPA, 1996) notes that if the calculated aquifer mixing zone depth is greater than the aquifer thickness, then the mixing zone depth should be set to equal to the aquifer thickness

It is noted that this method provides a DAF value that applies to leachate as it enters water bearing zones on the site, rather than at the discharge point, given the absence of any parameters that incorporate the overall catchment size or distance to the ultimate discharge point.

Major assumptions inherent to this method include:

- Dilution of the contaminant due to transport through the unsaturated soil zone is not included, the chemical in soil is assumed to be immediately adjacent to the water table; and
- Chemical degradation is also not included in this model at the soil contaminant source zone or in the resulting impacted groundwater, the calculations assume that the groundwater quality requirements must be achieved in the short term.

The above equations have been used to determine a DAF value for the site, based on site specific values for gradient, length of area parallel to groundwater flow direction and hydraulic conductivity. The calculated DAF value for the site using Equation 1 is presented in **Table 4**, along with the values used for determination.

Table 4:	Summary of DAF Calculation – USEPA 1996					
Parameter	Value	Reference				
i, hydraulic gradient (m/m)	3.1 x 10 ²	SKM 2006, highest gradient value reported for shallow wells				
da, aquifer thickness (m)	5	Estimated thickness of shallow water bearing zone based on SKM 2006 observations that groundwater in fill occurs as shallow as 1 m bgl, and that clay extends to approximately 6 m depth				
d, mixing zone depth (m)	5	Using equation 2 a value of 8.2 m was calculated for d. The aquifer thickness was therefore set as 5, adopting the recommendation in the method				

Parameter	Value	Reference				
I, infiltration rate (m/yr)	1.214	Bureau of Meteorology – Historical Annual Average Rainfall for Sydney (bom website on 28/07/10)- worst case scenario that 100% of rainfall infiltrates subsurface				
L, length of area of concern parallel to groundwater flow (m)	68	Maximum length of site running south-east				
K, aquifer conductivity (m/yr)	441.5	Based on maximum value of 1.4 x 10 ⁻⁵ ms ⁻¹ reported in SKM 2006 for shallow wells				
Calculated DAF	1.8	Using Equation 1 adopting the values listed above				

Given the low DAF calculated and the range of input data available for the site, DAFs were calculated for a range of scenarios, the results of which are summarised **Table 5** below. These values also provided as assessment of the sensitivity of the method.

-	Table 5:	Alternate DAF Calculations – adopting USEPA 1996					
Calculated DAF	% change	Scenarios					
3.76	109	Assuming only 30% of all rainfall on site infiltrates soil profile. Using a value of 0.3642 for I with all other values as per Table 4 .					
1.08	40	Assumes conductivity values are 1 order of magnitude below slug test calculated value, based on SKM 2006 report. Using a value of 44.15 m/yr for K with all other values as per Table 4 .					
1.01	44	Using a hydraulic gradient value of 4.41 x 10 ⁵ in the direction of groundwater flow. Based on the water levels reported in SKM 2006, at two shallow wells (17.57 m AHD at MW07S and 17.54 m AHD at MW03s) placed 68m apart					

Variation of the input values used in Equation 1 suggests that of all the parameters estimated for the site, the DAF calculated is most sensitive to the value adopted for infiltration.

The two main limitations of the USEPA (1996) method for the current application include:

- The method is intended to estimate the magnitude of dilution that occurs as soil leachate enters a water bearing zone on the site, rather than at the discharge point. Application of such a DAF to produce site specific criteria, is intended to result in GIL compliant groundwater migrating off site. Any parameters that incorporate the second stage of dilution that occurs between the site and the final discharge point, such as the overall catchment size or distance to the ultimate discharge point, are notably absent. For the current project this absence is considered to result in overly conservative DAF values. The site is located in an area where groundwater is unlikely to be utilised for primary contact, secondary contact or irrigation purposes. Under these circumstances, the environmental values of the catchment, or health impacts to future site users, will not be impacted if compliance with the GILs occurs at the site boundary or further along at the ultimate discharge point; and
- The limited sensitivity analysis conducted on the method focused on the parameters with the greatest potential for variation. The results indicated that the calculated DAF showed the greatest change as the value for the infiltration rate was altered (the DAF increased by 108% when the infiltration rate was reduced to 30% of the original input value, as opposed to only 40-44% change in the calculated DAF when values for hydraulic conductivity and hydraulic gradient were varied). As infiltration rate is governed by soil properties which are likely to be highly variable in shallow soils on the site, the approach of using literature review sourced values for input is also considered to be insufficiently robust for the current purpose.

Center for Research in Water Resources, University of Austin Texas (CRWR, 2003)

A report prepared by the Center for Research in Water Resources, University of Austin Texas (CRWR, 2003) describes the DAF calculation method used in the groundwater assessment component of the Texas Source Water Assessment Program. The Texas Source Water Assessment Program, was undertaken to determine the susceptibility of individual water sources to contamination.

The DAF applied to each water source was determined using the following series of equations:

$$DAF = DF \times AF$$

(Equation 3)

Where	DF AF	=	Dilution Factor, as calculated by Equation 4 below Attenuation Factor, as calculated by Equation 6 below				
	DF =	C _w /C _{soil}	={[$\rho_{b/}(\theta_{ws}+K_d\rho_b+H'\theta_{as})$] / LDF} x (L1/L2) (Equation 4)				
Where	C _w	=	Contaminant concentration in groundwater (g/cm ³)				
	C _{soil}	=	Contaminant concentration in soil (g/g-soil)				
	$ ho_{\scriptscriptstyle b}$	=	Soil bulk density (kg/L)				
	$ heta_{ws}$	=	the volumetric water content in the vadose zone (cm ³ -water/cm ³ -soil)				
	k _d	=	the soil water partitioning coefficient (cm ³ -water/g-soil)				
	H′	=	Henrys Law constant				
	$ heta_{as}$	=	the volumetric air content in the vadose zone (cm ³ -air/cm ³ -soil)				
	LDF	=	the Lateral Dilution Factor, as calculated by Equation 5 below				
	L1	=	thickness of affected soil				
	L2	=	Depth from the top of the affected soil to the groundwater table				
			δ)/(III()) (Equation 5)				

 $LDF = 1 + \{(U_{gw} \,\delta_{gw})/(I_fW_s)\}$

Where

(Equation 5)

U_{gw}	=	groundwater Darcy velocity (cm/year)
δ_{gw}	=	groundwater mixing zone thickness (m)
I_f	=	net infiltration rate (cm/year)
Ws	=	the lateral width of the affected vadose zone in the direction of groundwater flow(m)

 $AF = \exp[(L_{gw}/2a_x)^*(1 - \sqrt{1+4^*(D_ga_x/v_{coc})}] \times erf[W/4\sqrt{a_y}L_{gw})] \times erf[D/4\sqrt{a_z}L_{gw})]$ (Equation 6)

Where	L _{gw}	=	down gradient distance from the contamination source to the water supply well or discharge point (m)
	D_g	=	first order decay constant (day ⁻¹)
	V _{coc}	=	the contaminant retarded velocity (m/day)
	W	=	source zone width (m)
	D	=	source zone depth (m)
	a _{x,y,z}	=	the longitudinal(a_x), transverse (a_y) and vertical (a_z) groundwater dispersivities

Of all the DAF calculation methods considered, the CRWR method, requires the most input data, both in terms of volume and detail. Given the limited information on site hydrogeological conditions, and in particular on the fill layers, DAF calculations undertaken in this assessment have set the value of AF to 1. An AF value of 1, input into Equation 3, assumes that no attenuation processes are occurring within the water bearing zones, and any reduction in contaminant concentration that occurs is due to dilution only. This was considered to be suitably conservative for the current purpose.

The remaining values used as input into Equations 3 to 5 are presented in Table 6.

Table 6:	CRWR (2003) Method DAF Calculation Input Values
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Table 0.	0.0.0.0	5) Method DAP Calculation input values			
Parameter	Value	Reference			
i, hydraulic gradient (m/m)	3.1 x 10 ²	SKM 2006, highest gradient value reported for shallow wells			
$ ho_{b}$, soil bulk density	1.67	CRWR 2003, density value quoted for clayey soils			
θ_{ws}	0.16	CRWR 2003, value quoted for clayey soils			
θ_{as}	0.21	CRWR 2003, value quoted for clayey soils			
<i>k_d, the soil water partitioning coefficient (cm³-water/g-soil)</i>	Contaminant dependent	Used formula $k_d = f_{oc} \times K_{oc}$ Where f_{oc} value set to 0.002 g-carbon/ g-soil (CRWR, 2003), and K_{oc} values taken from (RAIS, 2010 ²) 1540 for naphthalene, 146 for benzene, 234 for toluene, 446 for ethylbenzene and 338 for xylene (all K_{oc} values in cm ³ - water/g soil)			
H', Henrys Law constant	Contaminant dependent	Values taken from (RAIS, 2010 ³) 0.018 for naphthalene, 0.322 for benzene, 0.271 for toluene, 0.322 for ethylbenzene and 0.212 for xylene (Henry's constant values unitless)			
L_1 , thickness of affected soil (m)	6	Estimated thickness of shallow fill and clay soils based on (CH2M Hill, 2007)			
L ₂ , Depth from top of affected soil to groundwater table (m)	1	Based on SKM 2006 observations that groundwater in fill occurs as shallow as 1 m bgl			
I_{f} , infiltration rate (m/yr)	1.214	Bureau of Meteorology – Historical Annual Average Rainfall for Sydney (bom website on 28/07/10)- worst case scenario that 100% of rainfall infiltrates subsurface			
$\delta g w$, groundwater mixing zone thickness (m)	5	Estimated thickness of shallow water bearing zone from SKM 2006 observations that groundwater in fill occurs as shallow as 1 m bgl, and clay extends to approximately 6 m depth			
<i>W_s, lateral width of affected vadose zone in direction of groundwater flow</i>	68	Maximum length of site running south-east			

Table 7 presents the spreadsheet calculations using the above values for the groundwater COPC identified in Section 3. Table 8 provides the DAF calculated for naphthalene using alternate input parameters.

The two main limitations of the CRWR (2003) method for the current application include:

- The model formulas require the input of values for detailed hydrogeological properties at the site, • e.g. volumetric water content, volumetric air contents, 'retarded contaminant velocity', which have not been determined for the site. While values can be, and have been, assumed for these properties based on literature reviews, this approach is considered to be insufficiently robust for the variable shallow water bearing zone under assessment; and
- The limited sensitivity analysis, focused on the parameters with the greatest potential for variation, indicated that the calculated DAF value showed the greatest change as the value for the fraction organic carbon content was increased. The value of this parameter is governed by soil properties which are likely to be highly variable in shallow soils on the site and potentially in soils located offsite. Insufficient data is available of catchment organic carbon levels to allow robust application for the current purpose.

² The Risk Assessment Information System (RAIS) website, Chemical Specific Parameters, <u>http://rais.ornl.gov/cgi-</u> <u>bin/tools/TOX_search?select=chem_spef</u> as on 29 July 2010 ³ The Risk Assessment Information System (RAIS) website, Chemical Specific Parameters, <u>http://rais.ornl.gov/cgi-</u>

bin/tools/TOX_search?select=chem_spef as on 29 July 2010

Contaminant	AF -	Ugw Cm/yr	δgw m	If Cm/yr	Ws m	LDF -	ρb Kg/L	Øws -	Foc g-carbon / g-soil	Koc Cm³-water /g soil	Kd Cm³-water /g soil	H′ -	θas -	L1 m	L2 m	DF -
naphthalene	1	620	5	121.4	68	1.376	1.67	0.16	0.002	1540	3.08	0.018	0.21	6	1	0.038
Benzene	1	620	5	121.4	68	1.376	1.67	0.16	0.002	146	0.292	0.322	0.21	6	1	0.283
Toluene	1	620	5	121.4	68	1.376	1.67	0.16	0.002	234	0.468	0.271	0.21	6	1	0.203
ethylbenzene	1	620	5	121.4	68	1.376	1.67	0.16	0.002	446	0.892	0.322	0.21	6	1	0.118
Xylenes	1	620	5	121.4	68	1.376	1.67	0.16	0.002	383	0.766	0.212	0.21	6	1	0.136

Table 7: CRWR Method DAF Calculations

 Table 8:
 CRWR method DAF Calculations for Naphthalene with Altered Input Values

Contaminant	AF -	Ugw Cm/yr	δgw m	If Cm/yr	Ws m	LDF -	ρb Kg/L	Ows -	Foc g- carbon / g- soil	Koc Cm ³ - water /g soil	Kd Cm ³ - water /g soil	H' -	θas -	L1 m	L2 m	DF -	% change
naphthalene	1	620	5	36.42 ¹	68	2.252	1.67	0.16	0.002	1540	3.08	0.018	0.21	6	1	0.0233	38.9
naphthalene	1	620	5	121.4	68	1.376	1.8 ²	0.16	0.002	1540	3.08	0.018	0.21	6	1	0.0382	0.2
naphthalene	1	620	5	121.4	68	1.376	1.67	0.16	0.02 ³	1540	30.8	0.018	0.21	6	1	0.0039	89.7
naphthalene	1	1370 ⁴	5	121.4	68	1.830	1.67	0.16	0.002	1540	3.08	0.018	0.21	6	1	0.0287	24.8

Notes:

1. Assuming only 30% of all rainfall on site infiltrates soil profile.

2. Assuming greater soil bulk density of 1.8 kg/L.

3. Assumes conductivity values are 1 order of magnitude below slug test calculated value, based on SKM 2006 report.

Using a hydraulic gradient value of 4.41 x 10⁵ in the direction of groundwater flow. Based on the water levels reported in SKM 2006, at two shallow wells (17.57 m AHD at MW07S and 17.54 m AHD at MW03s) placed 68m apart

Commonwealth Scientific and Industrial Research Organisation (CSIRO, 2009).

A report prepared by CSIRO (CSIRO, 2009)⁴ describes the National Environment Protection Council (NEPC) and Environment Protection and Heritage Council (EPHC) accepted and endorsed Australian method for deriving ecological investigation levels (EILs). The endorsed Australian method for deriving EILs is reported to be included as part of the revised National Environment Protection (Assessment of Site Contamination) Measure for public comment (CSIRO, 2009).

Appendix C of the CSIRO report (CSIRO, 2009) includes an endorsed method for deriving EILs that protect aquatic ecosystems. The method provides a means of calculating a dilution attenuation factor (DAF) for use in EIL calculations, which takes into account that groundwater catchments will most likely contain both contaminated and uncontaminated soils and pore water concentrations will not always equal groundwater concentrations of specific contaminants. The fraction of contaminated land to the total area of the groundwater/aquifer catchment is used to calculate the DAF as follows:

$DAF = 100 \div percentage of contaminated soil in catchment$ (Equation 7)

It is noted that in calculating a DAF the size of the whole catchment should be taken into consideration.

In calculating a DAF for the site using Equation 7, JBS has run two total catchment size scenarios to allow for the different discharge conditions that apply to shallow and deep groundwater from the site:

- Deep groundwater migrating south-east and discharging directly into Munni Street Creek and then Alexandra Canal; and
- Shallow perched water migrating south east off site and discharging into the piped stormwater channels along Railway Parade.

The DAF calculated for each catchment size scenario are provided in **Table 9** below. Adopting a conservative approach it was assumed that the areas of 'contaminated soil' comprised the entire site area.

	. ,					
Parameter	Alexandra Canal – deep groundwater	Reference	Railway Parade stormwater channel – shallow groundwater	Reference		
Total Area of Catchment (m ²)	16.6 x 10 ⁷	PPK 1999 ¹	117 600	PPK 1999 ¹		
Site Area (m ²)	7732	CH2M Hill 2007 ²	7732	CH2M Hill 2007 ²		
Ratio of 'Contaminated Soil' Area to Total Catchment Area	4.66 x 10 ⁻⁴	-	0.065748	-		
Percentage of Contaminated Soil in Catchment	0.05	-	7	-		
DAF	2000	-	16.6	-		

Table 9: CSIRO (2009) Method - Summary of DAF Calculations

Notes: 1 Catchment area estimated from PPK 1999 *"Cooks River Stormwater Management Plan*' Figure 2.4: Existing Stormwater Infrastructure, and Department of Lands Six Viewer website. Both figures included are as **Figure 2** and **Figure 3** respectively

While the CSIRO (2009) method appears to be the most simplistic of all the three methods considered, the basis of the method i.e. assessing the DAF at the point of discharge, rather than the site boundary, is considered to be the most appropriate for the project. The DAF value of 16.6 was calculated in a suitably conservative manner, i.e. assuming the nearest discharge point for shallow groundwater is the piped stormwater channel 50m south of the site on Railway Parade, and therefore reducing the catchment area size to less than 12 Ha. It is also noted that this calculated DAF is less than the USEPA (1996) default DAF value of 20 for site greater than 0.5 acres in area.

² CH2M Hill (December 2007) '*Remedial Action Plan*'.

⁴ The Australian Methodology to Derive Ecological Investigation levels in Contaminated Soils. CSIRO Land and Water Science Report 43/09, prepared for the National Environment Protection Council, (CSIRO, 2009).

7. Calculation of Site Specific Criteria for the Site

The three different approaches to DAF calculation have produced a range of results varying by four orders of magnitude. The CRWR method yields the most conservative results, which if adopted, would result in leachable concentrations criteria being only a small fraction of the corresponding GIL. The CSIRO method produces seemingly, the least conservative results. Consideration of the model assumptions and the anticipated context of application, has confirmed that the CSIRO (2009) method to be most suitable to be adopted for the site.

A value of 16.6 as determined by the CSIRO (2009) method, is the most appropriate DAF value for the site.

The DAF value of 16.6 has been used with the adopted GILs to derive of leachability criteria, as summarised in **Table 10**. The site specific leachability criteria assume the background concentrations of the contaminants of concern are zero, and that the leachable concentration in the treated material accounts for the total contaminant load at the discharge point. With the exception of heavy metals, this assumption is considered valid given the low concentrations of organic contaminants in wells located on the upper hydraulic gradient end of the site. While some heavy metal impact was noted in the groundwater migrating onto the site, this impact was considered to be representative of local conditions, and therefore it is considered adoption of the site specific values in **Table 10** for these contaminants would not adversely impact the environmental values at the point of discharge.

	Limit of Reporting	Aquatic Ecosystems ¹	Site Specific Criteria for assessment for leachable concentrations in soil ³
Arsenic (III/V)	0.1	2.3 ² / 4.5 ^{2, 5}	38.2 / 74.7
Cadmium	0.1	0.7 ¹	11.6
Chromium (III)	1	10	166
Chromium (VI)	0.1	4.4	73
Copper	0.1	1.3	21.6
Lead	0.1	4.4	73
Manganese	1	80 ²	1328
Mercury	0.05	0.1 ¹	1.76
Nickel	1	70	1162
Zinc	1	15	249
	VOLA	ATILE ORGANIC COMP	OUNDS
Benzene		500	8300
Toluene		180 ²	2988
Ethylbenzene		5 ²	83
Xylene (M+O+P)		625 ²	10 375
Styrene		1600	26560
Phenols		400	6640
	POLYCYC	LIC AROMATIC HYDRO	DCARBONS
Benzo(a)Pyrene	0.1	0.1 ²	1.7
Naphthalene	0.1	50	830
Phenanthrene	0.1	0.6 ²	10.3
Anthracene	0.1	0.01 ²	0.2
Fluoranthene	0.1	0.1 ²	1.7

Table 10: Leachate	Assessment Criteria	(all units in µg/L)
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¹ 95% protection levels (marine ecosystems) have been used. When these levels fail to protect key test species, the 99% protection levels were used - ANZECC/ARMCANZ (2000). The 99% protection levels have been adopted in line with recommendations in Section 8.3.7 of ANZECC/ARMCANZ 2000.

² Insufficient data to derive a reliable trigger value. In these instances, reference has been made to low reliability trigger levels contained in ANZECC/ARMCANZ (2000).

It is noted that TPH has not been included in **Table 10**, as the more prevalent indicator contaminants have been included.

It is concluded that the site specific criteria provided are over protective of the existing aquatic ecosystems of Alexandra Canal and can be used for future decision making for the site.

Prepared by,

Sumi Dorairaj Senior Environmental Consultant JBS Environmental Pty Ltd

Attachments:

(1) Limitations(2) Figures

Reviewed by,

M. Parke

Matthew Parkinson Principal - Contaminated Land JBS Environmental Pty Ltd

Attachment 1 - Limitations

This report has been prepared for use by Incoll Management Pty Ltd who commissioned the works in accordance with the project brief only and has been based in part on information obtained from other parties. The advice herein relates only to this project and all results conclusions and recommendations made should be reviewed by a competent person with experience in environmental investigations, before being used for any other purpose.

JBS Environmental Pty Ltd accepts no liability for use or interpretation by any person or body other than Incoll Management Pty Ltd or the appointed Site Auditor. This report should not be reproduced without prior approval by Incoll Management Pty Ltd, or amended in any way without prior approval by JBS Environmental Pty Ltd, and should not be relied upon by other parties, who should make their own enquires.

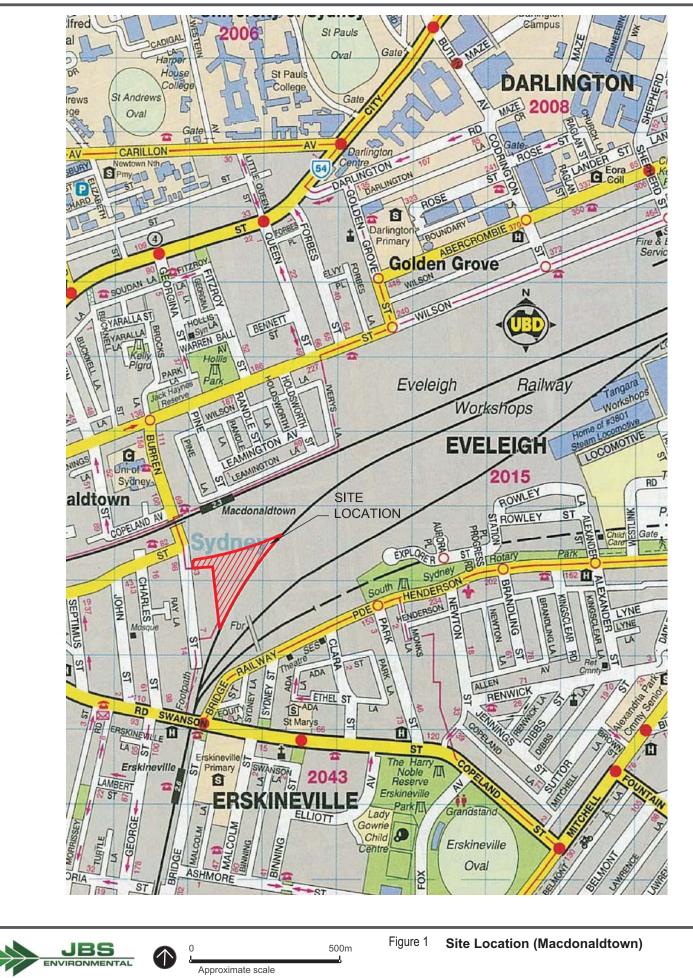
Sampling and chemical analysis of environmental media is based on appropriate guidance documents made and approved by the relevant regulatory authorities. Conclusions arising from the review and assessment of environmental data are based on the sampling and analysis considered appropriate based on the regulatory requirements and site history, not on sampling and analysis of all media at all locations for all potential contaminants.

Limited sampling and laboratory analyses were undertaken as part of the investigations, as described herein. Ground conditions between sampling locations may vary, and this should be considered when extrapolating between sampling points. Chemical analytes are based on the information detailed in the site history. Further chemicals or categories of chemicals may exist at the sites, which were not identified in the site history and which may not be expected at the site.

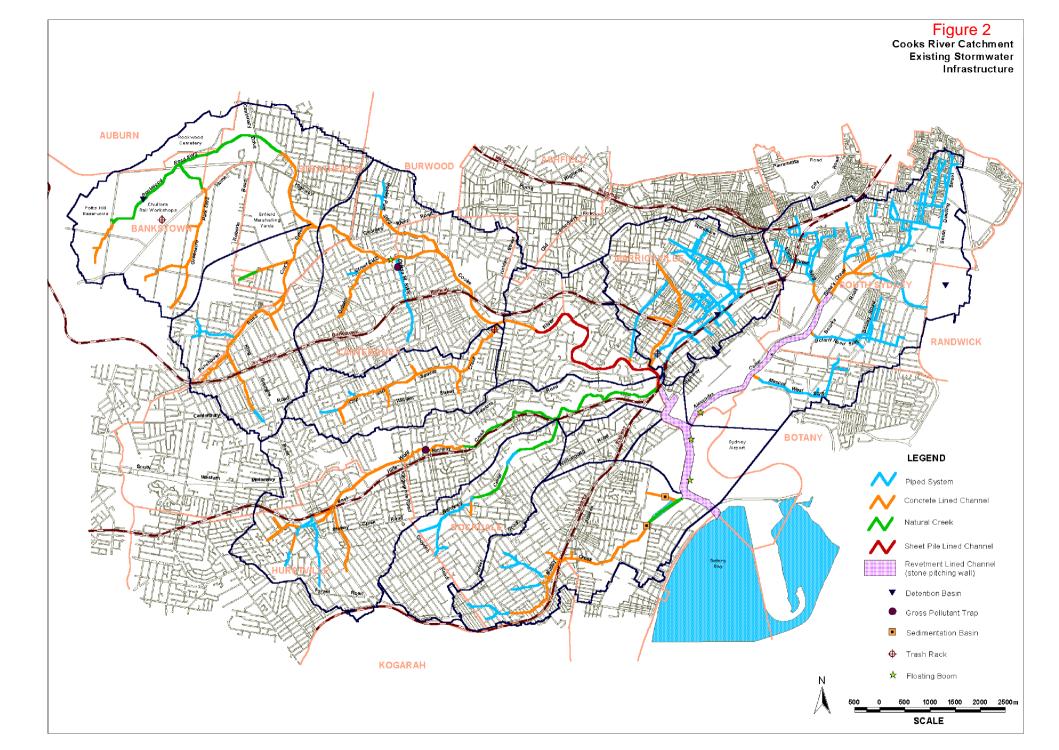
Changes to the subsurface conditions may occur subsequent to the investigations described herein, through natural processes or through the intentional or accidental addition of contaminants. The conclusions and recommendations reached in this report are based on the information obtained at the time of the investigations.

This report does not provide a complete assessment of the environmental status of the site, and it is limited to the scope defined herein. Should information become available regarding conditions at the site including previously unknown sources of contamination, JBS Environmental Pty Ltd reserves the right to review the report in the context of the additional information.

Attachment 2 Figures



CH2M Hill (2007) Note- All locations shown are approximate only



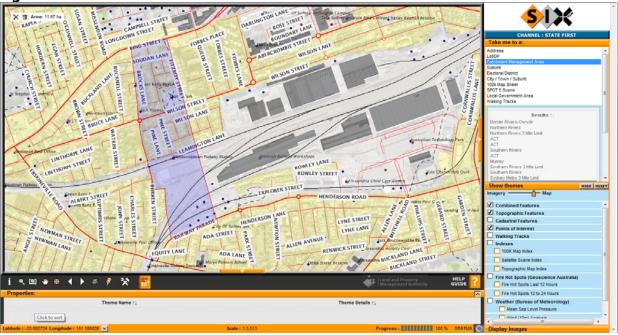


Figure 3: Extent of Shallow Groundwater Catchment Area

Purple shaded area showing extent of site catchment discharging into piped stormwater system leading to Munni Creek.



Appendix C

JBS Letter 'Results of Water Treatment Trial, former Macdonaldtown Gasworks, Burren Street Erskineville' JBS Environmental Pty Ltd (2010b)





JBS40913-15434 Rev 1

25 November 2010

John Dawson Project Director Incoll Management Pty Ltd Level 1, 73 Miller Street North Sydney NSW 2060 (Sent Via email: <u>jdawson@incoll.com.au</u>)

Groundwater Treatment Trial, Former Macdonaldtown Gasworks, Revision 1

Dear John,

1. Introduction and Objectives

JBS Environmental Pty Ltd (JBS) was engaged by Incoll Management Pty Ltd (Incoll). on behalf of Rail Corporation NSW Environment Projects Unit (RailCorp), to undertake works for the revision of the remedial strategy prepared for the former Macdonaldtown Gasworks site, This letter details the trial operation of a water treatment system on the site to assess the potential discharge locations for water collected during dewatering operations.

The former Macdonaldtown gasworks site has previously been the subject of environmental investigations which have identified a number of soil and groundwater contamination issues which require remediation and/or management in order to make the site suitable for ongoing commercial use. A remedial action plan (CH2M Hill 2007¹) for the site, has previously been prepared to document the procedures that will be undertaken to remediate and/or manage the identified issues at the site.

Excavation works proposed for the site include remedial works to remove contamination associated with naphthalene, benzo(a)pyrene (BaP) and Total Petroleum Hydrocarbons (TPH) within groundwater underlying the site. It is understood that proposed remedial excavations will extend to below the depth of groundwater underlying the site.

As a result of the contamination identified on the site, any groundwater collected during dewatering of the site for remediation is considered unsuitable to be disposed directly to stormwater. This 'dewater' will therefore require treatment prior to off-site disposal. JBS has undertaken a groundwater treatment trial, using a water treatment plant (WTP) transported to the site, to provide an indicative assessment of the 'treatability' of potential dewater from the site and an indicative assessment of treatment steps that will be required to manage the 'dewater'.

2. Methodology

Groundwater was extracted from existing monitoring wells MW37S, MW07S, MW06S and MW04S (**Figure 1**) from 05 July to 09 July 2010. The majority of water for the trial was pumped from MW04S and MW37S, which differed from the wells nominated for extraction in SAQP² for the works, as follows:

- SAQP nominated well MW03S was unable to be located at the time of the trial. The well was located in the vicinity of several large soil stockpiles, and it is uncertain whether this well remains viable;
- SAQP nominated well MW06S ran dry after extraction of approximately 40L on both 7 and 9 July 2010;
- SAQP nominated well MW07S ran dry after extraction of approximately 50L on both 7 and 9 July 2010;

¹ Remedial Action Plan, Former Macdonaltown Gasworks Site, Burren Street, Erskineville NSW 2007 (CH2M Hill, 2007)

² Sampling, Analysis and Quality Plan, Pre-Remedial Investigations, Former Macdonaltown Gasworks, JBS 2010

• SAQP nominated well MW31 was unable to be located at the time of the trial and it is likely this well has been destroyed.

Given the limited time available for completion of the trial, a total of only 4000L was pumped over the trial period from wells MW37S and MW04S. While not located within the source zones identified on site the groundwater pumped from MW37S and MW04S these wells is considered to be representative of impacted water.

Groundwater was extracted using bottom filling air operated in-well pumps installed to the base of the monitoring wells. Extracted groundwater was transferred to a new 9,000L storage poly tank. Approximately 4,000L of dewater was collected within the storage tanks during the pumping period.

The treatment system in use required a minimum of 8,000L of water to operate efficiently i.e. run for a sufficient length of time, and complete a meaningful trial i.e. allow for collection of sufficient samples. Consequently a further 4,000 L of water, sourced from on site fire hydrant was added to the 'dewater' in the influent tank on 9 July 2010. This approach was considered necessary because:

- Two of the four wells nominated for use in the trial were unable to be located and were presumed destroyed;
- The yield of groundwater in the two wells pumped in the vicinity of shallow groundwater plume was low, with a total of 4,000L pumped from both wells over the duration of the pumping trial.

The addition of water to influent tank for completion of the trial was considered not to impact the findings of the trial, given that the main objective of the trial was to assess the system's ability to remove the contaminants of concern from the 'dewater'. Provided the system is appropriate for the contaminants, treatment of water with higher concentrations will simply be a process of 'scaling up' the trialled components to the requirements of the remediation works. The findings of the trial are not intended to provide detailed design of the water treatment system, but rather to provide proposed remediation tenderers with sufficient information to anticipate the requirements of the remediation program.

The WTP used for the trial included the following components (listed in order of installation and water treatment):

- Oil / water separators;
- Multi-media (i.e. sand) filter (see Attachment 4); and
- Five 150L granular activated carbon (GAC) filters. Commercially available GAC was used in the filters. The amount of GAC used in the trial, equates to a GAC contact time during water treatment of 8 minutes.

Water flow through the WTP was controlled by the 92L/min flow restrictor provided to the oil-water separator. Effluent was discharged to an additional new sealed water storage tank. The water treatment trial was undertaken on the 9 July 2010.

Water sampling was undertaken at three intervals during the WTP operation (i.e. 'Run 1' collected 15 minutes after commencement, 'Run 2' collected 30mins after commencement and 'Run 3'collected 40 minutes after commencement). This series of runs was adopted to provide a 'snapshot' of water quality in the system between treatment components. The sampling strategy is listed in **Table 1** and depicted as a flow chart in **Figure 2**. Samples were collected from sampling 'taps' installed in pipe lines connecting the system components.

Sample ID	Sample Description
MW04S	Collected from undiluted groundwater held within storage tank prior to trial commencement.
Sample 1 – Run 1	Collected from water pumped out of influent tank, 15 minutes after trial commencement
Sample 1 – Run 2	Collected from water pumped out of influent tank, 30 minutes after trial commencement
Sample 1 – Run 3	Collected from water pumped out of influent tank, 40 minutes after trial commencement
Sample 2 – Run 1	Collected from water in WTP at outlet of oil / water separator at 15 minutes after trial commencement
Sample 2 – Run 2	Collected from water in WTP at outlet of oil / water separator at 30 minutes after trial commencement
Sample 2 – Run 3	Collected from water in WTP at outlet of oil / water separator at 40 minutes after trial commencement
Sample 3 – Run 1	Collected from water in WTP at outlet of air stripper at 15 minutes after trial commencement
Sample 3 – Run 2	Collected from water in WTP at outlet of air stripper at 30 minutes after trial commencement
Sample 3 – Run 3	Collected from water in WTP at outlet of air stripper 40 minutes after trial commencement
Sample 4 – Run 1	Collected from WTP effluent at 15 minutes. Additional treatment stages include sand filter, iron filter and GAC filter.
Sample 4 – Run 2	Collected from WTP effluent at 30 minutes. Additional treatment stages include sand filter, iron filter and GAC filter.
Sample 4 – Run 3	Collected from WTP effluent at 40 minutes. Additional treatment stages include sand filter, iron filter and GAC filter.

 Table 1:
 Summary of Water Treatment Trial Sampling

All samples were analysed for:

- Field parameters;
- Volatile organic compounds (VOCs);
- Total petroleum hydrocarbons (TPH);
- Polycyclic Aromatic Hydrocarbons (PAHs); and
- Heavy metals (including As, Cd, Cr, Cu, Pb, Hg, Ni and Zn) subsequent to field filtering.

3. Quality Assurance / Quality Control

A program of quality assurance / quality control (QA/QC) was undertaken with the sampling and analysis works conducted during the water treatment trial. QA/QC assessment included:

- Sampling and analysis of one inter-laboratory duplicate sample;
- Preparation and analysis of trip spike and trip blank samples;
- Conducting all sampling and sample preservation in accordance with JBS procedures;
- Use of NATA accredited laboratories for all analysis; and

• Assessment of laboratory QA/QC.

QA/QC has been assessed by reference to JBS standard quality protocols.

Based on the assessment of QA/QC, the environmental data generated during the water treatment trial are considered sufficiently representative to assess potential water treatment of groundwater underlying the site.

4. Results

Analytical results for the water treatment trial have been summarised in **Table 2**. Only COPCs which have been recorded above laboratory detection limits are summarised in **Table 2**. Laboratory result certificates are provided as **Attachment 3**.

From review of **Table 2**, observations during field works and the objectives of the water treatment trial the following points are observed:

- With the exception of arsenic, the concentrations of all heavy metals were significantly reduced by the system. Consistent reductions in these heavy metals concentrations were recorded after passing through each of the system components in all 3 'runs conducted, with the heavy metals concentrations in the final effluent samples from all runs (i.e. the Number 4 samples) less than the laboratory detection limits. ;
- Levels of arsenic and suspended solids increased slightly in effluent samples. GAC commonly has trace levels of arsenic contamination present which can become apparent in treated water. GAC also commonly releases fines, which can be recorded as suspended solids. This can be minimised by the selection of acid washed GAC filter for use in the operational WTP;
- Phase Separated Hydrocarbons (PSH) were not observed n the
- In Run 2, the majority of TPH C₁₀-C₃₆ removal was achieved by the oil-water separator. The significantly reduced level achieved at the 'Sample No. 2' location was further reduced to a concentrations less than the detection limits by the GAC filters;
- The results for Runs 1 and 3 indicate that the TPH C_{10} - C_{36} concentrations in the Sample 3 locations exceeded the Sample 2 values. As the plant operation and sampling methods used were consistent for the trial it is considered that these results are a reflection of the variable nature of groundwater under treatment, even after dilution. It is noted that regardless of this variation the concentrations of TPH C_{10} - C_{36} in the final effluent sample of both Runs 1 and 3 were less than the laboratory detection limits, and indicates the system provides an effective treatment process for these contaminants;
- The majority of PAH removal was achieved by the oil-water separator. This is most notable in naphthalene concentrations dropping by an order of magnitude between Sample 1 and Sample 2 locations for all three runs. Ultra-trace analysis was completed on the final effluent samples collected during the trial and in all three 'runs', all PAH concentrations were further reduced to concentrations less than the detection limits by the GAC filters;
- The majority of VOC removal was achieved by the GAC filters. GAC materials used in GAC filters are noted to have design lives, often measured as a 'breakthrough point'. Water treatment completed during dewatering will require to be cognisant of the mass of contaminants removed by the GAC and the relative absorption capacity; and
- Water used from the trial was removed from developed groundwater monitoring wells and had low levels of suspended solids. Levels of suspended solids from dewater generated during excavation works would be anticipated to have higher levels of dissolved solids.

Sample ID	рН					А	nalyte							
			VOCs		Т	эн				Heavy	Metals			
		Chloro- form	Bromo- dichloro- methane	Dibromo- chloro- methane	C ₆ -C ₉	C ₁₀ -C ₃₆	As	Cd	Cr	Cu	Pb	Hg	Ni	Zn
MW04S	Pumped groundwater holding tank	21	25	18	<10	6,200	<1	<0.1	<1	<1	<1	<0.5	1	110
Sample 1 – Run 1	Influent (15 mins)	14	8.9	2.7	<10	1,100	<1	<0.1	<1	<1	<1	<0.5	1	110
Sample 2 – Run 1	Post oil-water separator / pre sand filter (15 mins)	9.8	5.8	1.7	<10	170	<1	<0.1	<1	1	<1	<0.5	2	100
Sample 3 – Run 1	Post sand filter/ pre GAC (15 mins)	12	7.6	2.2	<10	3,210	<1	<0.1	<1	10	<1	<0.5	8	63
Sample 4 – Run 1	Post GAC Effluent (15 mins)	<1	<1	<1	<10	<250	16	<0.1	<1	<1	<1	<0.5	<1	1
Sample 1 – Run 2	Influent (30 mins)	12	2.1	2.1	<10	3,600	<1	<0.1	<1	3	<1	<0.5	7	170
Sample 2 – Run 2	Post oil-water separator / pre sand filter (30 mins)	11	6.6	2.0	<10	322	<1	<0.1	<1	4	<1	<0.5	9	91
Sample 3 – Run 2	Post sand filter/ pre GAC (30 mins)	9.7	6.3	2.0	<10	190	<1	<0.1	<1	7	<1	<0.5	8	48
Sample 4 – Run 2	Post GAC Effluent (30 mins)	<1	<1	<1	<10	<250	17	<0.1	<1	<1	<1	<0.5	<1	<1
Sample 1 – Run 3	Influent (40 mins)	9.6	5.9	1.8	<10	5,700	<1	<0.1	<1	<1	<1	<0.5	2	160
Sample 2 – Run 3	Post oil-water separator / pre sand filter (40 mins)	10	5.5	1.7	<10	<250	<1	<0.1	<1	<1	<1	<0.5	2	140
Sample 3 – Run 3	Post sand filter/ pre GAC (40 mins)	11	5.2	1.6	<10	1230	<1	<0.1	<1	4	1	<0.5	2	49
Sample 4 – Run 3	Post GAC Effluent (40 mins)	<1	<1	<1	<10	<250	24	<0.1	<1	<1	<1	<0.5	<1	<1

Table 2: Summary of Water Treatment Trial Results (all units in µg/L unless otherwise specified)

128 O'Riordan St Mascot NSW 2020

Table 2: Su	mmary of Water Treatme			(<u></u>					AHs					Analyte - PAHs										
Sample I D	Component or sampling stage (time after system commencement)	Naphthalene	Acenaphthylene	Acenaphthene	Fluorene	Phenanthrene	Anthracene	Fluoranthene	Pyrene	Benzo(a)anthracene	Chrysene	Benzo(b+k)fluoranthene	Benzo(a)pyrene	Indeno(1,2,3-c,d)pyrene	Dibenzo(a,h)anthracene	Benzo(g,h,i)perylene									
MW04S	Pumped groundwater holding tank	5.6	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1									
Sample 1 – Run 1	Influent (15 mins)	2.2	<0.1	0.7	0.5	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1									
Sample 2 – Run 1	Post oil-water separator / pre sand filter (15 mins)	0.3	<0.1	0.4	0.4	0.4	<0.1	0.2	0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1									
Sample 3 – Run 1	Post sand filter/ pre GAC (15 mins)	0.2	<0.1	0.3	0.2	0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1									
Sample 4 – Run 1	Post GAC (15 mins)	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.02	<0.01	<0.01	<0.01	<0.01									
Sample 1 – Run 2	Influent (30 mins)	3.4	<0.1	0.5	0.4	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1									
Sample 2 – Run 2	Post oil-water separator / pre sand filter (30 mins)	0.2	<0.1	0.5	0.5	1	0.2	0.7	0.4	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1									
Sample 3 – Run 2	Post sand filter/ pre GAC (30 mins)	0.2	<0.1	0.4	0.4	0.5	<0.1	<0.1	<0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1									
Sample 4 – Run 2	Post GAC (30 mins)	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.02	<0.01	<0.01	<0.01	<0.01									
Sample 1 – Run 3	Influent (45 mins)	3.8	<0.1	0.8	0.6	<0.1	<0.1	<0.1	<0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1									
Sample 2 – Run 3	Post oil-water separator / pre sand filter (40 mins)	0.2	<0.1	0.6	0.5	0.1	<0.1	0.1	<0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1									
Sample 3 – Run 3	Post sand filter/ pre GAC (40 mins)	0.2	<0.1	0.6	0.5	0.6	0.1	0.1	0.1	<0.1	<0.1	<0.2	<0.1	<0.1	<0.1	<0.1									
Sample 4 – Run 3	Post GAC (40 mins)	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.01	<0.02	<0.01	<0.01	<0.01	<0.01									

Table 2: Summary of Water Treatment Trial Results (all units in µg/L unless otherwise specified)

Overall it is noted that samples of the treatment system effluent (i.e. all 'Sample 4' results, collected post GAC filter) were reported to contain very low concentrations of all contaminants of concern at the site. Based on these results it appears that a water treatment plant could be used as part of the remediation process to enable discharge of collected groundwater as follows:

- To stormwater following receipt of relevant approvals from City of Sydney Council; or
- To sewer should a 'trade waste agreement' be entered into with Sydney Water.

Additionally, the results indicate that the treated effluent was of a suitable quality for reinjection into the subsurface. The feasibility of this option is however uncertain, given the required discharge rates during remediation are likely to exceed the infiltration potential of the clay soils underlying the site. This option would also require licensing by the NSW Office of Water.

If you wish to discuss any part of this letter further, then please free to contact Sumi Dorairaj on (02) 8338 1011. If you wish to discuss specifics of the WTP used during the works, or possible supply of water treatment components, Cameron Grant of Total Environmental Solutions (who supplied and operated the WTP during the trial) can be contacted on 0400 993 112.

Peer Review by:

Charlie Furr

Principal, Contaminated Land

JBS Environmental Pty Ltd

Prepared by:

Sumi Dorairaj Senior Environmental Consultant JBS Environmental Pty Ltd

Attachments:

(1) Limitations

(2) Figures(3) Laboratory Analysis Report and COC documentation(4) WTP media specifications

Attachment 1 – Limitations

This report has been prepared for use by the client who commissioned the works in accordance with the project brief only and has been based in part on information obtained from other parties. The advice herein relates only to this project and all results conclusions and recommendations made should be reviewed by a competent person with experience in environmental investigations, before being used for any other purpose.

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Sampling and chemical analysis of environmental media is based on appropriate guidance documents made and approved by the relevant regulatory authorities. Conclusions arising from the review and assessment of environmental data are based on the sampling and analysis considered appropriate based on the regulatory requirements and site history, not on sampling and analysis of all media at all locations for all potential contaminants.

Changes to the subsurface conditions may occur subsequent to the investigations described herein, through natural processes or through the intentional or accidental addition of contaminants. The conclusions and recommendations reached in this report are based on the information obtained at the time of the investigations.

This report does not provide a complete assessment of the environmental status of the site, and it is limited to the scope defined herein. Should information become available regarding conditions at the site including previously unknown sources of contamination, JBS Environmental Pty Ltd reserves the right to review the report in the context of the additional information.

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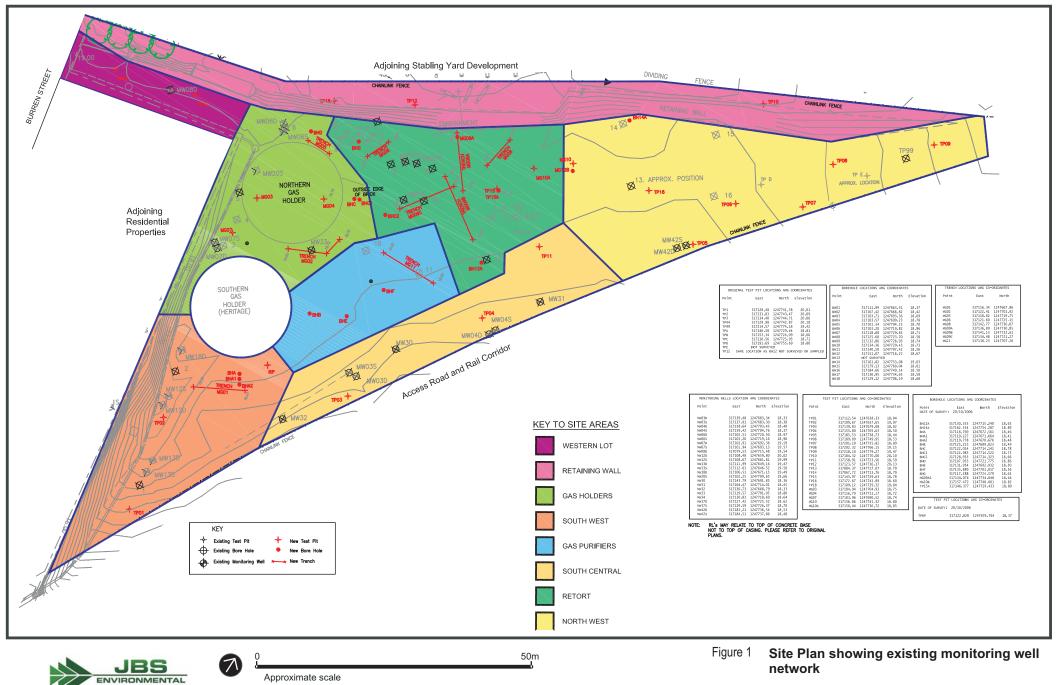
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Attachment 2 – Figures



CH2M Hill (2007) Note- All locations shown are approximate only

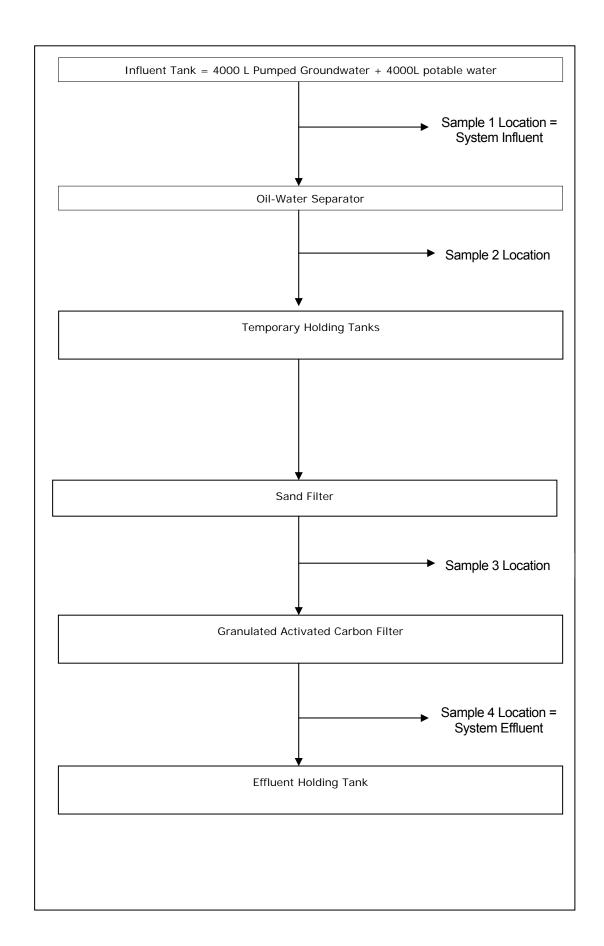


Figure 2: Water Treatment Trial - Sampling Strategy

Attachment 3 - Laboratory Results



Envirolab Services Pty Ltd ABN 37 112 535 645 12 Ashley St Chatswood NSW 2067 ph 02 9910 6200 fax 02 9910 6201 enquiries@envirolabservices.com.au www.envirolabservices.com.au

CERTIFICATE OF ANALYSIS 43277

Client: JBS Environmental P.O. Box 940 MASCOT NSW 1460

Attention: Tim Davis / Sumi Dorairaj

Sample log in details:

Your Reference: No. of samples: Date samples received: Date completed instructions received:

40913, Macdonaldtown

15 Waters 09/07/10 09/07/10

Analysis Details:

Please refer to the following pages for results, methodology summary and quality control data. Samples were analysed as received from the client. Results relate specifically to the samples as received. Results are reported on a dry weight basis for solids and on an as received basis for other matrices. *Please refer to the last page of this report for any comments relating to the results.*

Report Details:

 Date results requested by:
 20/07/10

 Date of Preliminary Report:
 Not issued

 Issue Date:
 23/07/10

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 Tests not covered by NATA are denoted with *.

Results Approved By:

Sandra Taylor Senior Organic Chemist

Kluign Morgen

Rhian Morgan Metals Supervisor

Jacinta Hurst Laboratory Manager

Envirolab Reference: 432 Revision No: R 0

43277 R 01

Nick Sarlamis Inorganics Supervisor



Page 1 of 23

VOCs in water						
Our Reference:	UNITS	43277-1	43277-2	43277-3	43277-4	43277-5
Your Reference		Sample 1 -	Sample 2 -	Sample 3 -	Sample 4 -	Sample 1 -
		Run 1	Run 1	Run 1	Run 1	Run 2
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water	Water	Water
Date extracted	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/2010
Date analysed	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/2010
Dichlorodifluoromethane	µg/L	<10	<10	<10	<10	<10
Chloromethane	µg/L	<10	<10	<10	<10	<10
Vinyl Chloride	µg/L	<10	<10	<10	<10	<10
Bromomethane	µg/L	<10	<10	<10	<10	<10
Chloroethane	µg/L	<10	<10	<10	<10	<10
Trichlorofluoromethane	µg/L	<10	<10	<10	<10	<10
1,1-Dichloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Trans-1,2-dichloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1-dichloroethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Cis-1,2-dichloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Bromochloromethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Chloroform	µg/L	14	9.8	12	<1.0	12
2,2-dichloropropane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichloroethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,1-trichloroethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1-dichloropropene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Cyclohexane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Carbon tetrachloride	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Dibromomethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichloropropane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Trichloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Bromodichloromethane	µg/L	8.9	5.8	7.6	<1.0	7.1
trans-1,3-dichloropropene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
cis-1,3-dichloropropene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,2-trichloroethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Toluene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,3-dichloropropane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Dibromochloromethane	μg/L	2.7	1.7	2.2	<1.0	2.1
1,2-dibromoethane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Tetrachloroethene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,1,2-tetrachloroethane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Chlorobenzene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Ethylbenzene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Bromoform	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
m+p-xylene	μg/L	<2.0	<2.0	<2.0	<2.0	<2.0
Styrene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,2,2-tetrachloroethane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
.,.,_,_ totaonioi ootilailo	۳9'۲	1.5				



VOCs in water						
Our Reference:	UNITS	43277-1	43277-2	43277-3	43277-4	43277-5
Your Reference		Sample 1 - Run 1	Sample 2 - Run 1	Sample 3 - Run 1	Sample 4 - Run 1	Sample 1 - Run 2
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water	Water	Water
o-xylene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,3-trichloropropane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Isopropylbenzene		<1.0	<1.0	<1.0	<1.0	<1.0
Bromobenzene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
	μg/L	-	-	-	-	-
n-propyl benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
2-chlorotoluene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
4-chlorotoluene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,3,5-trimethyl benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Tert-butyl benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,4-trimethyl benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,3-dichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Sec-butyl benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,4-dichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
4-isopropyl toluene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
n-butyl benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dibromo-3-chloropropane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,4-trichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Hexachlorobutadiene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,3-trichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Surrogate Dibromofluoromethane	%	96	104	99	108	104
Surrogate toluene-d8	%	108	109	115	110	108
Surrogate 4-BFB	%	105	103	106	105	106

VOCs in water						
Our Reference:	UNITS	43277-6	43277-7	43277-8	43277-9	43277-10
Your Reference		Sample 2 -	Sample 3 -	Sample 4 -	Sample 1 -	Sample 2 -
		Run 2	Run 2	Run 2	Run 3	Run 3
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water	Water	Water
Date extracted	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/2010
Date analysed	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/2010
Dichlorodifluoromethane	μg/L	<10	<10	<10	<10	<10
Chloromethane	μg/L	<10	<10	<10	<10	<10
Vinyl Chloride	μg/L	<10	<10	<10	<10	<10
Bromomethane	µg/L	<10	<10	<10	<10	<10
Chloroethane	µg/L	<10	<10	<10	<10	<10
Trichlorofluoromethane	µg/L	<10	<10	<10	<10	<10
1,1-Dichloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Trans-1,2-dichloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1-dichloroethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Cis-1,2-dichloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Bromochloromethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Chloroform	µg/L	11	9.7	<1.0	9.6	10
2,2-dichloropropane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichloroethane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,1-trichloroethane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1-dichloropropene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Cyclohexane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Carbon tetrachloride	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Benzene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Dibromomethane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichloropropane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Trichloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Bromodichloromethane	µg/L	6.6	6.3	<1.0	5.9	5.5
trans-1,3-dichloropropene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
cis-1,3-dichloropropene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,2-trichloroethane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Toluene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,3-dichloropropane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Dibromochloromethane	μg/L	2.0	2.0	<1.0	1.8	1.7
1,2-dibromoethane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Tetrachloroethene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,1,2-tetrachloroethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Chlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Ethylbenzene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Bromoform	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
m+p-xylene	μg/L	<2.0	<2.0	<2.0	<2.0	<2.0
Styrene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,2,2-tetrachloroethane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1, 1, ∠, ∠ ⁻ leu do noi 0eu idite	ру/L	\$1.0	\$1.0	\$1.0	51.0	51.0

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VOCs in water						
Our Reference:	UNITS	43277-6	43277-7	43277-8	43277-9	43277-10
Your Reference		Sample 2 - Run 2	Sample 3 - Run 2	Sample 4 - Run 2	Sample 1 - Run 3	Sample 2 - Run 3
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water	Water	Water
o-xylene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,3-trichloropropane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Isopropylbenzene		<1.0	<1.0	<1.0	<1.0	<1.0
Bromobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
	µg/L	-	-	-	-	_
n-propyl benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
2-chlorotoluene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
4-chlorotoluene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,3,5-trimethyl benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Tert-butyl benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,4-trimethyl benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,3-dichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Sec-butyl benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,4-dichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
4-isopropyl toluene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
n-butyl benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dibromo-3-chloropropane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,4-trichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Hexachlorobutadiene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,3-trichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Surrogate Dibromofluoromethane	%	98	97	102	100	101
Surrogate toluene-d8	%	108	112	109	108	92
Surrogate 4-BFB	%	106	105	106	106	105

VOCs in water				
Our Reference:	UNITS	43277-11	43277-12	43277-13
Your Reference		Sample 3 - Run 3	Sample 4 - Run 3	MW04s
Date Sampled		9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water
Date extracted	-	13/07/2010	13/07/2010	13/07/2010
Date analysed	-	13/07/2010	13/07/2010	13/07/2010
Dichlorodifluoromethane	µg/L	<10	<10	<10
Chloromethane	µg/L	<10	<10	<10
Vinyl Chloride	µg/L	<10	<10	<10
Bromomethane	µg/L	<10	<10	<10
Chloroethane	µg/L	<10	<10	<10
Trichlorofluoromethane	µg/L	<10	<10	<10
1,1-Dichloroethene	µg/L	<1.0	<1.0	<1.0
Trans-1,2-dichloroethene	µg/L	<1.0	<1.0	<1.0
1,1-dichloroethane	µg/L	<1.0	<1.0	<1.0
Cis-1,2-dichloroethene	µg/L	<1.0	<1.0	<1.0
Bromochloromethane	µg/L	<1.0	<1.0	<1.0
Chloroform	µg/L	11	<1.0	21
2,2-dichloropropane	µg/L	<1.0	<1.0	<1.0
1,2-dichloroethane	µg/L	<1.0	<1.0	<1.0
1,1,1-trichloroethane	µg/L	<1.0	<1.0	<1.0
1,1-dichloropropene	µg/L	<1.0	<1.0	<1.0
Cyclohexane	µg/L	<1.0	<1.0	<1.0
Carbon tetrachloride	µg/L	<1.0	<1.0	<1.0
Benzene	µg/L	<1.0	<1.0	<1.0
Dibromomethane	µg/L	<1.0	<1.0	<1.0
1,2-dichloropropane	µg/L	<1.0	<1.0	<1.0
Trichloroethene	µg/L	<1.0	<1.0	<1.0
Bromodichloromethane	µg/L	5.2	<1.0	25
trans-1,3-dichloropropene	µg/L	<1.0	<1.0	<1.0
cis-1,3-dichloropropene	µg/L	<1.0	<1.0	<1.0
1,1,2-trichloroethane	µg/L	<1.0	<1.0	<1.0
Toluene	µg/L	<1.0	<1.0	<1.0
1,3-dichloropropane	µg/L	<1.0	<1.0	<1.0
Dibromochloromethane	µg/L	1.6	<1.0	18
1,2-dibromoethane	µg/L	<1.0	<1.0	<1.0
Tetrachloroethene	µg/L	<1.0	<1.0	<1.0
1,1,1,2-tetrachloroethane	µg/L	<1.0	<1.0	<1.0
Chlorobenzene	µg/L	<1.0	<1.0	<1.0
Ethylbenzene	µg/L	<1.0	<1.0	<1.0
Bromoform	µg/L	<1.0	<1.0	<1.0
m+p-xylene	µg/L	<2.0	<2.0	<2.0
Styrene	µg/L	<1.0	<1.0	<1.0
1,1,2,2-tetrachloroethane	µg/L	<1.0	<1.0	<1.0



VOCs in water				
Our Reference:	UNITS	43277-11	43277-12	43277-13
Your Reference		Sample 3 -	Sample 4 -	MW04s
		Run 3	Run 3	
Date Sampled		9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water
o-xylene	μg/L	<1.0	<1.0	<1.0
1,2,3-trichloropropane	µg/L	<1.0	<1.0	<1.0
Isopropylbenzene	µg/L	<1.0	<1.0	<1.0
Bromobenzene	μg/L	<1.0	<1.0	<1.0
n-propyl benzene	μg/L	<1.0	<1.0	<1.0
2-chlorotoluene	μg/L	<1.0	<1.0	<1.0
4-chlorotoluene	µg/L	<1.0	<1.0	<1.0
1,3,5-trimethyl benzene	µg/L	<1.0	<1.0	<1.0
Tert-butyl benzene	μg/L	<1.0	<1.0	<1.0
1,2,4-trimethyl benzene	μg/L	<1.0	<1.0	<1.0
1,3-dichlorobenzene	μg/L	<1.0	<1.0	<1.0
Sec-butyl benzene	μg/L	<1.0	<1.0	<1.0
1,4-dichlorobenzene	μg/L	<1.0	<1.0	<1.0
4-isopropyl toluene	μg/L	<1.0	<1.0	<1.0
1,2-dichlorobenzene	µg/L	<1.0	<1.0	<1.0
n-butyl benzene	μg/L	<1.0	<1.0	<1.0
1,2-dibromo-3-chloropropane	μg/L	<1.0	<1.0	<1.0
1,2,4-trichlorobenzene	µg/L	<1.0	<1.0	<1.0
Hexachlorobutadiene	μg/L	<1.0	<1.0	<1.0
1,2,3-trichlorobenzene	µg/L	<1.0	<1.0	<1.0
Surrogate Dibromofluoromethane	%	95	100	101
Surrogate toluene-d8	%	77	76	108
Surrogate 4-BFB	%	105	105	104

Client Reference: 40913, Ma

40913, Macdonaldtown

vTPH & BTEX in Water						
Our Reference:	UNITS	43277-1	43277-2	43277-3	43277-4	43277-
Your Reference		Sample 1 - Run 1	Sample 2 - Run 1	Sample 3 - Run 1	Sample 4 -	Sample Run 2
Date Sampled		9/07/2010	9/07/2010	9/07/2010	Run 1 9/07/2010	9/07/20
Type of sample		Water	Water	Water	Water	Water
		13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/20
Date extracted	-					
Date analysed	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/20
TPH C6 - C9	µg/L	<10	<10	<10	<10	<10
Surrogate Dibromofluoromethane	%	96	104	99	108	104
Surrogate toluene-d8	%	108	109	115	110	108
Surrogate 4-BFB	%	105	103	106	105	106
vTPH & BTEX in Water						
Our Reference:	UNITS	43277-6	43277-7	43277-8	43277-9	43277-
Your Reference		Sample 2 -	Sample 3 -	Sample 4 -	Sample 1 -	Sample
Data Samplad		Run 2 9/07/2010	Run 2 9/07/2010	Run 2 9/07/2010	Run 3 9/07/2010	Run 3 9/07/20
Date Sampled Type of sample		Water	Water	Water	Water	9/07/20 Wate
Date extracted	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/20
Date analysed	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/20
TPH C6 - C9	µg/L	<10	<10	<10	<10	<10
Surrogate Dibromofluoromethane	%	98	97	102	100	101
Surrogate toluene-d8	%	108	112	109	108	92
Surrogate 4-BFB	%	106	105	106	106	105
vTPH & BTEX in Water						
Our Reference:	UNITS	43277-11	43277-12	43277-13	43277-14	43277-
Your Reference		Sample 3 - Run 3	Sample 4 - Run 3	MW04s	Trip Blank	Trip Spi
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/20
Type of sample		Water	Water	Water	Water	Wate
Date extracted	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/20
Date analysed	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/20
TPH C6 - C9	µg/L	<10	<10	<10	[NA]	[NA]
Benzene	µg/L	[NA]	[NA]	[NA]	<1.0	108%
Toluene	µg/L	[NA]	[NA]	[NA]	<1.0	129%
Ethylbenzene	μg/L	[NA]	[NA]	[NA]	<1.0	97%
m+p-xylene	μg/L	[NA]	[NA]	[NA]	<2.0	95%
o-xylene	μg/L	[NA]	[NA]	[NA]	<1.0	96%
Surrogate Dibromofluoromethane	%	95	100	100	96	91
Surrogate toluene-d8	%	77	76	107	109	124
Surrogate 4-BFB	%	105	105	104	105	103

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sTPH in Water (C10-C36)						
Our Reference:	UNITS	43277-1	43277-2	43277-3	43277-4	43277-5
Your Reference		Sample 1 -	Sample 2 -	Sample 3 -	Sample 4 -	Sample 1
		Run 1	Run 1	Run 1	Run 1	Run 2
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water	Water	Water
Date extracted	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/201
Date analysed	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/201
TPH C10 - C14	µg/L	<50	<50	<50	<50	<50
TPH C15 - C28	µg/L	1,100	170	810	<100	2,100
TPH C29 - C36	µg/L	<100	<100	2,400	<100	1,500
Surrogate o-Terphenyl	%	96	87	#	100	80
sTPH in Water (C10-C36)			10077 7		10077 0	
Our Reference:	UNITS	43277-6	43277-7	43277-8	43277-9	43277-1
Your Reference		Sample 2 - Run 2	Sample 3 - Run 2	Sample 4 - Run 2	Sample 1 - Run 3	Sample 2 Run 3
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/201
Type of sample		Water	Water	Water	Water	Water
Date extracted	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/201
Date analysed	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/201
TPH C10 - C14	μg/L	52	<50	<50	<50	<50
TPH C15 - C28	μg/L	270	190	<100	3,500	<100
TPH C29 - C36	µg/L	<100	<100	<100	2,200	<100
Surrogate o-Terphenyl	%	92	77	86	99	86
sTPH in Water (C10-C36)						

sTPH in Water (C10-C36)				
Our Reference:	UNITS	43277-11	43277-12	43277-13
Your Reference		Sample 3 -	Sample 4 -	MW04s
		Run 3	Run 3	
Date Sampled		9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water
Date extracted	-	13/07/2010	13/07/2010	13/07/2010
Date analysed	-	13/07/2010	13/07/2010	13/07/2010
TPH C10 - C14	µg/L	<50	<50	<50
TPH C15 - C28	µg/L	410	<100	3,800
TPH C29 - C36	µg/L	820	<100	2,800
Surrogate o-Terphenyl	%	96	93	73

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PAHs in Water - Low Level						
Our Reference:	UNITS	43277-1	43277-2	43277-3	43277-5	43277-6
Your Reference		Sample 1 -	Sample 2 -	Sample 3 -	Sample 1 -	Sample 2 -
		Run 1	Run 1	Run 1	Run 2	Run 2
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water	Water	Water
Date extracted	-	13/7/2010	13/7/2010	13/7/2010	13/7/2010	13/7/2010
Date analysed	-	13/7/2010	13/7/2010	13/7/2010	13/7/2010	13/7/2010
Naphthalene	µg/L	2.2	0.3	0.2	3.4	0.2
Acenaphthylene	µg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Acenaphthene	µg/L	0.7	0.4	0.3	0.5	0.5
Fluorene	µg/L	0.5	0.4	0.2	0.4	0.5
Phenanthrene	µg/L	<0.1	0.4	0.1	<0.1	1.0
Anthracene	µg/L	<0.1	<0.1	<0.1	<0.1	0.2
Fluoranthene	µg/L	<0.1	0.2	<0.1	<0.1	0.7
Pyrene	µg/L	<0.1	0.1	<0.1	<0.1	0.4
Benzo(a)anthracene	µg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Chrysene	µg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Benzo(b+k)fluoranthene	µg/L	<0.2	<0.2	<0.2	<0.2	<0.2
Benzo(a)pyrene	µg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Indeno(1,2,3-c,d)pyrene	μg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Dibenzo(a,h)anthracene	μg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Benzo(g,h,i)perylene	μg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Surrogate p-Terphenyl-d14	%	129	111	129	123	99



PAHs in Water - Low Level						
Our Reference:	UNITS	43277-7	43277-9	43277-10	43277-11	43277-13
Your Reference		Sample 3 -	Sample 1 -	Sample 2 -	Sample 3 -	MW04s
		Run 2	Run 3	Run 3	Run 3	0/07/0040
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water	Water	Water
Date extracted	-	13/7/2010	13/7/2010	13/7/2010	13/7/2010	13/7/2010
Date analysed	-	13/7/2010	13/7/2010	13/7/2010	13/7/2010	13/7/2010
Naphthalene	µg/L	0.2	3.8	0.2	0.2	5.6
Acenaphthylene	µg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Acenaphthene	µg/L	0.4	0.8	0.6	0.6	<0.1
Fluorene	µg/L	0.4	0.6	0.5	0.5	<0.1
Phenanthrene	µg/L	0.5	<0.1	0.1	0.6	<0.1
Anthracene	µg/L	<0.1	<0.1	<0.1	0.1	<0.1
Fluoranthene	µg/L	<0.1	<0.1	0.1	0.1	0.2
Pyrene	µg/L	<0.1	<0.1	<0.1	0.1	0.2
Benzo(a)anthracene	µg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Chrysene	µg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Benzo(b+k)fluoranthene	µg/L	<0.2	<0.2	<0.2	<0.2	<0.2
Benzo(a)pyrene	μg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Indeno(1,2,3-c,d)pyrene	μg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Dibenzo(a,h)anthracene	μg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Benzo(g,h,i)perylene	μg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Surrogate p-Terphenyl-d14	%	84	127	111	127	87



PAH low level in Water				
Our Reference:	UNITS	43277-4	43277-8	43277-12
Your Reference		Sample 4 -	Sample 4 -	Sample 4 -
		Run 1	Run 2	Run 3
Date Sampled		9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water
Date extracted	-	22/07/2010	22/07/2010	22/07/2010
Date analysed	-	23/7/2010	23/7/2010	23/7/2010
Naphthalene	µg/L	<0.01	<0.01	<0.01
Acenaphthylene	µg/L	<0.01	<0.01	<0.01
Acenaphthene	µg/L	<0.01	<0.01	<0.01
Fluorene	µg/L	<0.01	<0.01	<0.01
Phenanthrene	µg/L	<0.01	<0.01	<0.01
Anthracene	µg/L	<0.01	<0.01	<0.01
Fluoranthene	µg/L	<0.01	<0.01	<0.01
Pyrene	µg/L	<0.01	<0.01	<0.01
Benz(a)anthracene	µg/L	<0.01	<0.01	<0.01
Chrysene	µg/L	<0.01	<0.01	<0.01
Benzo(b)&(k)fluoranthene	µg/L	<0.02	<0.02	<0.02
Benzo(a)pyrene	µg/L	<0.01	<0.01	<0.01
Indeno(1,2,3-cd)pyrene	µg/L	<0.01	<0.01	<0.01
Dibenz(ah)anthracene	µg/L	<0.01	<0.01	<0.01
Benzo(ghi)perylene	µg/L	<0.01	<0.01	<0.01

Envirolab Reference: 43277 Revision No: R 01 ACCREDITED FOR TECHNICAL COMPETENCE

Total Phenolics in Water						
Our Reference:	UNITS	43277-1	43277-2	43277-3	43277-4	43277-5
Your Reference		Sample 1 -	Sample 2 -	Sample 3 -	Sample 4 -	Sample 1 -
		Run 1	Run 1	Run 1	Run 1	Run 2
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water	Water	Water
Date extracted	-	14/7/2010	14/7/2010	14/7/2010	14/7/2010	14/7/2010
Date analysed	-	14/7/2010	14/7/2010	14/7/2010	14/7/2010	14/7/2010
Total Phenolics (as Phenol)	mg/L	<0.050	<0.050	<0.050	0.080	<0.050
Total Phenolics in Water						
Our Reference:	UNITS	43277-6	43277-7	43277-8	43277-9	43277-10
Your Reference		Sample 2 -	Sample 3 -	Sample 4 -	Sample 1 -	Sample 2 -
		Run 2	Run 2	Run 2	Run 3	Run 3
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water	Water	Water
Date extracted	-	14/7/2010	14/7/2010	14/7/2010	14/7/2010	14/7/2010
Date analysed	-	14/7/2010	14/7/2010	14/7/2010	14/7/2010	14/7/2010
Total Phenolics (as Phenol)	mg/L	<0.050	<0.050	<0.050	0.12	<0.050
					7	
Total Phenolics in Water		10077.11	10077 10	10077 10		
Our Reference:	UNITS	43277-11	43277-12	43277-13		
Your Reference		Sample 3 - Run 3	Sample 4 - Run 3	MW04s		
Date Sampled		9/07/2010	9/07/2010	9/07/2010		
Type of sample		Water	Water	Water		
Date extracted		14/7/2010	14/7/2010	14/7/2010	=	
Date analysed	_	14/7/2010	14/7/2010	14/7/2010		
-						

<0.050

< 0.050

< 0.050

mg/L

Total Phenolics (as Phenol)



HM in water - dissolved						
Our Reference:	UNITS	43277-1	43277-2	43277-3	43277-4	43277-5
Your Reference		Sample 1 -	Sample 2 -	Sample 3 -	Sample 4 -	Sample 1 -
- /		Run 1	Run 1	Run 1	Run 1	Run 2
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water	Water	Water
Date prepared	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/2010
Date analysed	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/2010
Arsenic-Dissolved	µg/L	<1	<1	<1	16	<1
Cadmium-Dissolved	μg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Chromium-Dissolved	µg/L	<1	<1	<1	<1	<1
Copper-Dissolved	µg/L	<1	1	10	<1	3
Lead-Dissolved	µg/L	<1	<1	<1	<1	<1
Mercury-Dissolved	µg/L	<0.5	<0.5	<0.5	<0.5	<0.5
Nickel-Dissolved	µg/L	1	2	8	<1	7
Zinc-Dissolved	μg/L	110	100	63	1	170
HM in water - dissolved						
Our Reference:	UNITS	43277-6	43277-7	43277-8	43277-9	43277-10
Your Reference		Sample 2 -	Sample 3 -	Sample 4 -	Sample 1 -	Sample 2
		Run 2	Run 2	Run 2	Run 3	Run 3
Date Sampled		9/07/2010	9/07/2010	9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water	Water	Water
Date prepared	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/201
Date analysed	-	13/07/2010	13/07/2010	13/07/2010	13/07/2010	13/07/201
Arsenic-Dissolved	µg/L	<1	<1	17	<1	<1
Cadmium-Dissolved	µg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Chromium-Dissolved	µg/L	<1	<1	<1	<1	<1
Copper-Dissolved	µg/L	4	7	<1	<1	<1
Lead-Dissolved	µg/L	<1	<1	<1	<1	<1
Mercury-Dissolved	μg/L	<0.5	<0.5	<0.5	<0.5	<0.5
Nickel-Dissolved	μg/L	9	8	<1	2	2
					100	

91

48

<1

160

Zinc-Dissolved

µg/L

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HM in water - dissolved				
Our Reference:	UNITS	43277-11	43277-12	43277-13
Your Reference		Sample 3 -	Sample 4 -	MW04s
		Run 3	Run 3	
Date Sampled		9/07/2010	9/07/2010	9/07/2010
Type of sample		Water	Water	Water
Date prepared	-	13/07/2010	13/07/2010	13/07/2010
Date analysed	-	13/07/2010	13/07/2010	13/07/2010
Arsenic-Dissolved	µg/L	<1	24	<1
Cadmium-Dissolved	µg/L	<0.1	<0.1	<0.1
Chromium-Dissolved	µg/L	<1	<1	<1
Copper-Dissolved	µg/L	4	<1	9
Lead-Dissolved	µg/L	1	<1	<1
Mercury-Dissolved	µg/L	<0.5	<0.5	<0.5
Nickel-Dissolved	µg/L	2	<1	3
Zinc-Dissolved	µg/L	49	<1	45

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Method ID	Methodology Summary
GC.13	Water samples are analysed directly by purge and trap GC-MS.
GC.16	Soil samples are extracted with methanol and spiked into water prior to analysing by purge and trap GC-MS. Water samples are analysed directly by purge and trap GC-MS.
GC.3	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-FID.
GC.12 subset	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-MS.
Ext-020	Analysis subcontracted to Australian Government - National Measurement Institute. NATA Accreditation No: 198
LAB.30	Total Phenolics - determined colorimetrically following disitillation.
Metals.22 ICP-MS	Determination of various metals by ICP-MS.
Metals.21 CV-AAS	Determination of Mercury by Cold Vapour AAS.



QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
VOCs in water						Base II Duplicate II %RPD		Recovery
Date extracted	-			13/07/2 010	[NT]	[NT]	LCS-W1	13/07/2010
Date analysed	-			13/07/2 010	[NT]	[NT]	LCS-W1	13/07/2010
Dichlorodifluoromethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Chloromethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Vinyl Chloride	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Bromomethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Chloroethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Trichlorofluoromethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
1,1-Dichloroethene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Trans-1,2-dichloroethen e	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,1-dichloroethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	112%
Cis-1,2-dichloroethene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Bromochloromethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Chloroform	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	110%
2,2-dichloropropane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,2-dichloroethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	105%
1,1,1-trichloroethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	105%
1,1-dichloropropene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Cyclohexane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Carbon tetrachloride	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Dibromomethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,2-dichloropropane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Trichloroethene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	114%
Bromodichloromethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	111%
trans-1,3-dichloropropen e	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
cis-1,3-dichloropropene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,1,2-trichloroethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Toluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,3-dichloropropane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Dibromochloromethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	113%
1,2-dibromoethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Tetrachloroethene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	110%
1,1,1,2-tetrachloroethan e	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Chlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Ethylbenzene	μg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Bromoform	μg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
m+p-xylene	μg/L	2	GC.13	<2.0	[NT]	[NT]	[NR]	[NR]
Styrene	μg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]

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40913. Macdonaldtown

Client Reference: 40913, Macdonaldtown											
QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery			
VOCs in water						Base II Duplicate II %RPD					
1,1,2,2-tetrachloroethan e	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
o-xylene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
1,2,3-trichloropropane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
Isopropylbenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
Bromobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
n-propyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
2-chlorotoluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
4-chlorotoluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
1,3,5-trimethyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
Tert-butyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
1,2,4-trimethyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
1,3-dichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
Sec-butyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
1,4-dichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
4-isopropyl toluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
1,2-dichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
n-butyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
1,2-dibromo-3-chloropro pane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
1,2,4-trichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
Hexachlorobutadiene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
1,2,3-trichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]			
<i>Surrogate</i> Dibromofluoromethane	%		GC.13	102	[NT]	[NT]	LCS-W1	114%			
Surrogate toluene-d8	%		GC.13	101	[NT]	[NT]	LCS-W1	107%			
Surrogate 4-BFB	%		GC.13	101	[NT]	[NT]	LCS-W1	100%			

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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
vTPH & BTEX in Water						Base II Duplicate II %RPD		-
Date extracted	-		T	13/07/2 010	[NT]	[NT]	LCS-W1	13/07/2010
Date analysed	-			13/07/2 010	[NT]	[NT]	LCS-W1	13/07/2010
TPH C6 - C9	µg/L	10	GC.16	<10	[NT]	[NT]	LCS-W1	113%
Benzene	µg/L	1	GC.16	<1.0	[NT]	[NT]	[NR]	[NR]
Toluene	µg/L	1	GC.16	<1.0	[NT]	[NT]	[NR]	[NR]
Ethylbenzene	µg/L	1	GC.16	<1.0	[NT]	[NT]	[NR]	[NR]
m+p-xylene	µg/L	2	GC.16	<2.0	[NT]	[NT]	[NR]	[NR]
o-xylene	µg/L	1	GC.16	<1.0	[NT]	[NT]	[NR]	[NR]
<i>Surrogate</i> Dibromofluoromethane	%		GC.16	102	[NT]	[NT]	LCS-W1	110%
Surrogate toluene-d8	%		GC.16	101	[NT]	[NT]	LCS-W1	104%
Surrogate 4-BFB	%		GC.16	101	[NT]	[NT]	LCS-W1	101%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
sTPH in Water (C10-C36)						Base II Duplicate II %RPD		
Date extracted	-			13/07/2 010	[NT]	[NT]	LCS-W2	13/07/2010
Date analysed	-			13/07/2 010	[NT]	[NT]	LCS-W2	13/07/2010
TPH C10 - C14	µg/L	50	GC.3	<50	[NT]	[NT]	LCS-W2	69%
TPH C15 - C28	µg/L	100	GC.3	<100	[NT]	[NT]	LCS-W2	83%
TPH C29 - C36	µg/L	100	GC.3	<100	[NT]	[NT]	LCS-W2	86%
Surrogate o-Terphenyl	%		GC.3	94	[NT]	[NT]	LCS-W2	100%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
PAHs in Water - Low Level						Base II Duplicate II %RPD		
Date extracted	-			13/7/20 10	[NT]	[NT]	LCS-W2	13/7/2010
Date analysed	-			13/7/20 10	[NT]	[NT]	LCS-W2	13/7/2010
Naphthalene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-W2	96%
Acenaphthylene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Acenaphthene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Fluorene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-W2	112%
Phenanthrene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-W2	104%
Anthracene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]

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			ent Referen		0913, Macdon		Craike Craff	Spike %
QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Recovery
PAHs in Water - Low Level						Base II Duplicate II %RPD		
Fluoranthene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-W2	104%
Pyrene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-W2	110%
Benzo(a)anthracene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Chrysene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-W2	117%
Benzo(b+k)fluoranthene	µg/L	0.2	GC.12 subset	<0.2	[NT]	[NT]	[NR]	[NR]
Benzo(a)pyrene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-W2	131%
Indeno(1,2,3-c,d)pyrene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Dibenzo(a,h)anthracene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Benzo(g,h,i)perylene	µg/L	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Surrogate p-Terphenyl-d ₁₄	%		GC.12 subset	75	[NT]	[NT]	LCS-W2	77%
QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike %
			METHOD	Dianix	Duplique onin	Duplicate recould	opike onin	Recovery
PAH low level in Water						Base II Duplicate II %RPD		
Date extracted	-			22/07/2 010	[NT]	[NT]	LCS-W1	22/7/2010
Date analysed	-			23/07/2 010	[NT]	[NT]	LCS-W1	23/7/2010
Naphthalene	µg/L	0.01	Ext-020	<0.01	[NT]	[NT]	LCS-W1	90%
Acenaphthylene	µg/L	0.01	Ext-020	<0.01	[NT]	[NT]	[NR]	[NR]
Acenaphthene	µg/L	0.01	Ext-020	<0.01	[NT]	[NT]	[NR]	[NR]
Fluorene	µg/L	0.01	Ext-020	<0.01	[NT]	[NT]	LCS-W1	92%
Phenanthrene	µg/L	0.01	Ext-020	<0.01	[NT]	[NT]	LCS-W1	89%
Anthracene	µg/L	0.01	Ext-020	<0.01	[NT]	[NT]	[NR]	[NR]
Fluoranthene	µg/L	0.01	Ext-020	<0.01	[NT]	[NT]	[NR]	[NR]
Pyrene	µg/L	0.01	Ext-020	<0.01	[NT]	[NT]	[NR]	[NR]
				1				

µg/L

µg/L

µg/L

µg/L

µg/L

µg/L

µg/L

Benz(a)anthracene

Chrysene Benzo(b)&(k)fluoranthen

е

Benzo(a)pyrene

Indeno(1,2,3-cd)pyrene

Dibenz(ah)anthracene

Benzo(ghi)perylene

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0.01

0.01

0.02

0.01

0.01

0.01

0.01

Ext-020

Ext-020

Ext-020

Ext-020

Ext-020

Ext-020

Ext-020

<0.01

<0.01

< 0.02

<0.01

<0.01

<0.01

<0.01

[NT]



[NR]

LCS-W1

[NR]

LCS-W1

[NR]

LCS-W1

[NR]

[NR]

91%

[NR]

79%

[NR]

89%

[NR]

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike %
								Recovery
Total Phenolics in Water						Base II Duplicate II %RPD		
Date extracted	-			14/7/20 10	43277-1	14/7/2010 14/7/2010	43277-1	14/7/2010
Date analysed	-			14/7/20 10	43277-1	14/7/2010 14/7/2010	43277-1	14/7/2010
Total Phenolics (as Phenol)	mg/L	0.05	LAB.30	<0.050	43277-1	<0.050 <0.050	43277-1	87%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
HM in water - dissolved						Base II Duplicate II %RPD		,
Date prepared	-			13/07/2 010	43277-1	13/07/2010 13/07/201	0 LCS-W1	13/07/2010
Date analysed	-			13/07/2 010	43277-1	13/07/2010 13/07/201	0 LCS-W1	13/07/2010
Arsenic-Dissolved	µg/L	1	Metals.22 ICP-MS	<1	43277-1	<1 <1	LCS-W1	106%
Cadmium-Dissolved	µg/L	0.1	Metals.22 ICP-MS	<0.1	43277-1	<0.1 <0.1	LCS-W1	109%
Chromium-Dissolved	µg/L	1	Metals.22 ICP-MS	<1	43277-1	<1 <1	LCS-W1	96%
Copper-Dissolved	µg/L	1	Metals.22 ICP-MS	<1	43277-1	<1 <1	LCS-W1	93%
Lead-Dissolved	µg/L	1	Metals.22 ICP-MS	<1	43277-1	<1 <1	LCS-W1	101%
Mercury-Dissolved	µg/L	0.5	Metals.21 CV-AAS	<0.5	43277-1	<0.5 <0.5	LCS-W1	117%
Nickel-Dissolved	µg/L	1	Metals.22 ICP-MS	<1	43277-1	1 1 RPD: 0	LCS-W1	93%
Zinc-Dissolved	µg/L	1	Metals.22 ICP-MS	<1	43277-1	110 110 RPD: 0	LCS-W1	99%
QUALITY CONTROL Total Phenolics in Water	UNITS	S	Dup. Sm#	Base +	Duplicate Duplicate + %RPD	Spike Sm#	Spike % Recovery	
Date extracted	-		43277-11	14/7/2	2010 14/7/2010	43277-2	14/7/2010	
Date analysed	-		43277-11	14/7/2	2010 14/7/2010	43277-2	14/7/2010	
Total Phenolics (as Phenol	l) mg/L	-	43277-11	<0.	.050 <0.050	43277-2	98%	
QUALITY CONTROL	UNITS	S	Dup. Sm#		Duplicate	Spike Sm#	Spike % Recovery	
HM in water - dissolved				Base +	Duplicate + %RPD)		
Date prepared	-		43277-11	13/07/2	010 13/07/2010	43277-2	13/07/2010	
Date analysed	-		43277-11	13/07/2	2010 13/07/2010	43277-2	13/07/2010	
Arsenic-Dissolved	µg/L	.	43277-11		<1 <1	43277-2	116%	
Cadmium-Dissolved	μg/L		43277-11		<0.1 <0.1	43277-2	120%	
Chromium-Dissolved µg/L			43277-11		<1 <1	43277-2	102%	
Copper-Dissolved µg/L		.	43277-11	4	4 RPD: 0	43277-2	95%	
Lead-Dissolved	µg/L	.	43277-11		1 <1	43277-2	102%	
Mercury-Dissolved	µg/L	.	43277-11	.	<0.5 <0.5	43277-2	115%	

Envirolab Reference: 43277 Revision No: R 01 ACCREDITED FOR TECHNICAL COMPETENCE

Client Reference: 40913, Macdonaldtown												
QUALITY CONTROL HM in water - dissolved	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery							
Nickel-Dissolved	µg/L	43277-11	2 2 RPD: 0	43277-2	96%							
Zinc-Dissolved	µg/L	43277-11	49 49 RPD: 0	43277-2	105%							

Envirolab Reference: 43277 Revision No: R 01



Report Comments:

Total Petroleum Hydrocarbons in water (semivol):# Percent recovery is not possible to report as the high concentration of analytes in the sample/s have caused interference.

PAH's in water analysed by NMI. Report No.RN806180.

Asbestos was analysed by Approved Identifier: Not applicable for this job

Asbestos was authorised by Approved Signatory: Not applicable for this job

INS: Insufficient sample for this test NT: Not tested PQL: Practical Quantitation Limit <: Less than >: Greater than **RPD: Relative Percent Difference** NA: Test not required LCS: Laboratory Control Sample NR: Not requested

Quality Control Definitions

Blank: This is the component of the analytical signal which is not derived from the sample but from reagents, glassware etc, can be determined by processing solvents and reagents in exactly the same manner as for samples. Duplicate: This is the complete duplicate analysis of a sample from the process batch. If possible, the sample selected should be one where the analyte concentration is easily measurable.

Matrix Spike: A portion of the sample is spiked with a known concentration of target analyte. The purpose of the matrix spike is to monitor the performance of the analytical method used and to determine whether matrix interferences exist.

LCS (Laboratory Control Sample): This comprises either a standard reference material or a control matrix (such as a blank sand or water) fortified with analytes representative of the analyte class. It is simply a check sample.

Surrogate Spike: Surrogates are known additions to each sample, blank, matrix spike and LCS in a batch, of compounds which are similar to the analyte of interest, however are not expected to be found in real samples.

Laboratory Acceptance Criteria:

Duplicate sample and matrix spike recoveries may not be reported on smaller jobs, however, were analysed at a frequency to meet or exceed NEPM requirements. All samples are tested in batches of 20. The duplicate sample RPD and matrix spike recoveries for the sample batch were within laboratory acceptance criteria.

Duplicates: <5xPQL - any RPD is acceptable; >5xPQL - 0-50% RPD is acceptable.

Matrix Spikes and LCS: Generally 70-130% for inorganics/metals; 60-140% for organics and 10-140% for

SVOC and speciated phenols is acceptable. Surrogates: 60-140% is acceptable for general organics and 10-140% for

Envirolab Reference: Revision No:

43277 R 01



CHAIN OF CUSTODY - Client

ENVIROLAB SERVICES



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Client: J&						Client Project Name and Number:								Envirolab Services 12 Ashley St, Chatswood, NSW, 2067				
Project Mgr: S. DORAIRAG					Macdonaldtown PO No.: 40913													
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Attachment 4 – WTP Filter Specifications



Automatic Multi Media Water Filters

For Removal Of Suspended Matter

MODELS: FMA-A, FMA-F, FMA-F40 & FMA-F50

The multi layered media filter has advantages over conventional filters :

- * Greater Dirt Holding Capacity
- * Higher Flow Rates
- * Greater ability to handle surges
- * Reduced Backwash Rates

SPECIFICATIONS:

All units automatically backwash and rinse on a periodic basis in order to cleanse and reclassify the filter media. This is normally accomplished by an adjustable timer.

Sediment filters incur an increase pressure loss between inlet and outlet as filtered particles collect in the bed.

OPTIONS:

- * Backwash Only
- * Flush after Backwash
- * Choice of Timers or Signal Start
- * Custom Design
- * Pressure Gauges
- * Pressure Limiting & Relief Valves



FMA 550F40

FMA - A & FMA - F PIPE SIZE: 25mm - Maximum Flowrate 110L/min

FLOW RATES						DIMENSIONS					
	Model	Difficult ⁽¹⁾ L/min	Normal ⁽²⁾ L/min	Maximum ⁽³⁾ L/min	Backwash ⁽⁴⁾ L/min	Bed Area m ²	Cylinder Dia. mm	Height mm	Shipping Weight		
	FMA 200A	12	16	28	19	0.032	200	1200	50		
	FMA 250A	19	24	44	29	0.050	250	1200	65		
	FMA 300F or	·A-P 29	36	65	43	0.074	300	1200	80		
	FMA 325F or	·A-P 33	42	75	50	0.086	325	1500	86		
	FMA 350F or	·A-P 39	49	87	58	0.100	350	1500	90		
	FMA 400F or	·A-P 49	62	110	74	0.126	400	1200	160		
	FMA 550F	97	110	110	120 ⁽⁵⁾	0.250	550	1400	200		

FMA - F40 PIPE SIZE: 40mm - Maximum Flowrate 254L/min

						-			
FLOW RATES/PRESSURE DROPS					DIMENSIONS				
Model	Difficult ⁽¹⁾ L/min/kPa	Normal ⁽²⁾ L/min/kPa	Maximum ⁽³⁾ L/min/kPa	Backwash ⁽⁴⁾ L/min	Bed Area m ²	Cylinder Dia. mm	Height mm	Shipping Weight	
FMA 400F40	49/20	62/30	110/60	74	0.12	400	1900	150	
FMA 550F40	97/50	122/60	219/140	146	0.25	550	1400	200	
FMA 600F40	113/55	142/80	254/185	170	0.29	600	2200	350	
FMA 750F40	175/100	220/140	254/185	182	0.45	750	2600	700	

FMA - F50 PIPE SIZE: 50mm - Maximum Flowrate 580L/min

FMA 900F50	257/50	323/120	577/200	380	0.66	900	2600	1000
		020/220	0, = 00	000	0.00	500	2000	2000

(1) Removal of compressible solids eg. Floc carryover at feed concentrations to 300gm/L.

Reduces pressure loss and increases period between backwashes. Based on 390L/min/m².

(2) Removal of incompressible suspended solids above 10 microns at feed concentrations to 300gm/L.

Based on 490L/min/m².

(3) Not recommended for continuous use. Based on 875L/min/m².

(4) Based on 585L/min/m².

(5) Special limit minimum pressure 450 kPa.

* SPECIFICATIONS SUBJECT TO CHANGE WITHOUT NOTICE. IF ANY DIMENSIONS ARE CRITICAL CONFIRM WHEN PLACING ORDER.



Appendix D Sydney Water Trade Waste Acceptance Criteria Brochure

Sydney WATER

Industrial customers

Acceptance standards and charging rates for 2010-11

Sydney Water accepts trade wastewater to the sewer, if it meets certain acceptance standards.

What are acceptance standards?

Acceptance standards are generally limits to the concentration of substances, in composite samples of trade wastewater discharge. For substances that pose a particular health and safety risk, acceptance standards also apply to the concentration of substances in a discrete sample of trade wastewater discharge.

These substances are highlighted in **bold** print in the tables in this fact sheet.

The acceptance standards for domestic substances are listed in Table 1, and for non-domestic in Table 2. Table 1 also lists the value of the domestic equivalent concentrations, which we deduct when calculating charges.

The Independent Pricing and Regulatory Tribunal (IPART) determined that all trade waste fees and charges will be adjusted from 1 July (under Determination No.1, 2008)

Testing

Customers must make sure substances specified in trade waste agreements or permits are only analysed in laboratories registered by the National Association of Testing Authorities (NATA), for the class of test(s) or specific test(s).

The approved analytical methods may be downloaded from <u>sydneywater.com.au</u>

What are they based on?

Acceptance standards are based on:

- safe levels of substances that may otherwise pose a health risk to workers in and around the sewerage system
- safe levels of substances to protect public health
- pollution reduction targets and discharge licence conditions set by the Department of Environment, Climate Change and Water NSW (DECCW NSW)
- the need to protect our assets and treatment processes
- the capability of the sewerage system to transport 'domestic substances', ie suspended solids, grease and BOD
- concentrations obtainable by using proven pre-treatment technology (provision is made to trial new technology)
- quality specifications for biosolids and reuse water
- reuse considerations, including the need to provide wastewater that does not interfere with reuse treatment processes, or limit reuse opportunities
- national acceptance criteria published as *Guidelines for Sewerage Systems, Acceptance of Trade Waste (Industrial Waste)*, (ARMCANZ & ANZECC, November 1994).



•	· ·			
Substance	Acceptance standard (mg/L)	Domestic equivalent (mg/L)	Note	Charging rate (\$/kg)
Suspended solids	600	200		0.862
BOD5 - primary treatment		230	1	0.120 + {0.0178 x (BOD/600)}
BOD5 - secondary treatment		230	1	0.678 + {0.0178 x (BOD/600)}
Soluble BOD	100	Not applicable	15	0.120 + {0.0178 x (BOD/600)}
Grease – primary treatment	110	50	2	1.214
Grease – secondary treatment	200	50	2	1.214
Ammonia	100	35	3, 5	2.014
Nitrogen	150	50	4	0.170
Phosphorus	50	10	4	1.347
Sulphate	2000	50		0.133 x (SO ₄ /2000)
Total dissolved solids (ocean systems, no discharge limitation)	10000	450	12	0.0058
Total dissolved solids (inland and ocean systems with limitation)	500	450	12	0.0058
Total dissolved solids (inland and ocean systems with advanced treatment to remove TDS)	10000	450	12, 16	0.173 x fraction of average dry weather flow treated

Table 1: Acceptance standards, domestic equivalents and charging rates for domestic substances

Trade waste requirements

- Sydney Water will determine standards for colour and interference with ultra violet disinfection on a system-specific basis.
- There must be no fibrous material in the trade wastewater, if we believe it could obstruct or block the sewerage system.
- Non-faecal gross solids must have a maximum linear dimension of less than 20 mm, a maximum cross section of 6 mm and must have a quiescent settling velocity of less than 3 m/hr.
- Sydney Water will negotiate radioactive material activity rates for sewer discharge on a site-specific basis.

 The Manager, Business Customer Delivery will determine acceptance standards for substances other than those listed in Tables 1 and 2. Sydney Water does not accept substances (or mixtures of substances) that cannot mix with water.

Provisional standards

Where we determine that an additional substance should be included on our list of acceptance standards, the new acceptance standard will be declared provisional.

The substance will be provisional for six months. During this time, the customer must test for the substance, but no charges will be levied.

There are currently no provisional standards.



Substance	Acceptance standard (mg/L)	Note	Charging rate (\$/kg)
Acetaldehyde	5	5	13.502
Acetone	400	5	0.127
Aluminium	100		0.677
Arsenic	1		67.574
Barium	5		13.502
Boron	100		0.677
Bromine	5	5	13.502
Cadmium	1		67.574
Chlorinated phenolics	0.05	6	1351.675
Chlorine	10	5	6.756
Chromium	3	7	22.281
Cobalt	5		13.502
Copper	5		13.502
Cyanide	1	5, 8	67.574
Fluoride	20	4	3.346
Formaldehyde	30	5	2.234
General pesticides (excludes OC and OP)	0.1	9	675.811
Herbicides and defoliants	0.1	-	675.811
Iron	50		1.344
Lead	2		33.750
Lithium (specified systems only)	10	10	6.756
Manganese	10		6.756
Mercaptans	1		67.574
Mercury	0.03		2230.222
Methyl Ethyl Ketone	100	5	0.677
Molybdenum	100		0.677
Nickel	3		22.281
Organoarsenic compounds	0.1		675.811
pH	7-10 units	1	
Petroleum hydrocarbons (flammable)	10	5, 11, 14,18	6.756
- Benzene	0.1	5, 18	
- Toluene	0.5	5, 18	
- Ethylbenzene	1	5, 18	
- Xylene	1	5, 18	
Phenolic compounds (non-chlorinated)	1	0,10	67.574
Polynuclear aromatic hydrocarbons	5		13.502
Propionaldehyde	5	5	13.502
Selenium	5	0	13.502
Silver	5		13.502
Sulphide	5	5	13.502
Sulphite	50	5	1.344
Temperature	38°C	1	1.044
Thiosulphate	300	I	0.243
Tin	10		6.756
Uranium	10		
	10	5 10 17	6.756
Volatile halocarbons		5, 13, 17	67.574
- Chloroform	0.1	5, 17	
- Perchloroethylene	0.3	5, 17	
- Trichloroethylene	0.1	5, 17	10 500
Zinc	5		13.502

Table 2 Acceptance standards and charging rates for non-domestic substances



Notes to acceptance standards

- 1. Sydney Water will introduce acceptance standards for a substance on a sub-system specific basis as determined by:
 - how much the receiving system can transport and treat
 - how corroded the sub-system is
 - how sewage treatment products will be used.
- 2. Discrete oil, fat or grease must not be discharged.
- 3. Where ammonia is present with other nitrogenous compounds, the amount of nitrogen in the ammonia is deducted from the total nitrogen as measured by Total Kjeldahl Nitrogen, before calculating the charge for nitrogen.
- 4. Fluoride, phosphorus and nitrogen limits don't apply where the customer's sewerage system is connected to a sewage treatment plant that discharges to the ocean.
- 5. Acceptance standards also apply to concentrations of ammonia, benzene, bromine, chlorine, cyanide, formaldehyde, petroleum hydrocarbons, sulphide and volatile halocarbons in discrete samples.
- We will determine acceptance standards for individual chlorinated phenolics on a catchment basis, following pollution reduction targets set by the DECCW NSW for the sewage treatment plant effluent. The concentration limit is a guide only and we may set lower limits for individual chlorinated phenolic compounds.
- 7. We do not allow discharge from comfort air conditioning cooling towers and evaporative condensers using products containing hexavalent chromium (chromate) or organometallic algicides, if the blow down (or 'bleed-off') is connected to the sewer. Comfort cooling towers are defined as cooling towers dedicated to heating, ventilation, air-conditioning or refrigeration systems.
- 8. Cyanide is defined as labile cyanide amenable to alkaline chlorination. This includes free cyanide as well as those complex cyanides that are particularly dissociable, almost wholly, or in a large degree, and therefore potentially toxic in low concentrations.

- We will not consent to any discharge of organochlorine pesticides (including chlordane, dieldrin and heptachlor), or organophosphorus pesticides (including chlorpyrifos, diazinon and malathion) into the sewerage system.
- 10. The limit for lithium applies only to the Rouse Hill sewage catchment.
- 11. Where flammable and/or explosive substances may be present, the customer must demonstrate to us that there is no possibility of explosions, or fires in the sewerage system. We will discuss limits and charges with individual customers, before a trade waste agreement is negotiated. The flammability of the discharge must never exceed five per cent of the Lower Explosive Limit (LEL) of hexane at 25 °C. In some cases a customer may be required to install an LEL meter.
- 12. We will determine acceptance standards for total dissolved solids on a catchment-specific basis. A limit of 500 mg/L may apply to customers discharging to an inland sewage treatment plant or to a sewage treatment plant that is part of a designated reuse system. Acceptance standards will only apply to those customers discharging in excess of 100kg/d of total dissolved solids (TDS) or greater than one per cent of the total catchment TDS load (whichever is the lesser).
- 13. Analysis of volatile halocarbons must at a minimum include methylene chloride, chloroform, trichloroethylene and perchloroethylene.
- 14. This substance is made up of several substances including benzene, toluene, ethylbenzene, (m+p)-xylene and o-xylene.
- 15. As at 1 July 2010, the limit for soluble BOD applies only to the Smithfield sewage and SPS 67 catchments, due to corrosion.
- 16. This is a guide only. Exact \$/kg rates are determined on a system-specific basis.
- 17. Charges will apply for total volatile halocarbons
- 18. Charges will apply for total petroleum hydrocarbons (flammable)

Want to know more?

Visit sydneywater.com.au



Appendix E DECC Immobilisation Technical Note 1



You are here: <u>Home</u> > <u>Waste and resource recovery</u> > <u>Regulating waste in NSW</u> > <u>Waste immobilisation</u> > <u>Technical notes</u> > Note 1

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Immobilisation Technical Note 1

Process Equipment for Treatment of Contaminated Soil and Sludge Waste

In the context of hazardous waste treatment under the POEO Act, it is imperative that the responsible person uses proper process plant and equipment competently to conduct the treatment. This approach would help assure that the favourable treatment results achieved at the laboratory or pilot scale trials can be repeated consistently at the full scale treatment and high treatment standards are maintained at all times.

Industry may practise either or both of the following methods in the immobilisation treatment of contaminated soil and sludge waste:

- 1. <u>Chemical fixation</u>: Chemical reagents are used to convert the target contaminants contained in the waste to a chemically stable form(s) suitable for landfill disposal.
- 2. <u>Stabilisation/solidification</u>: Cement and/or pozzolans reagents are used to transform the waste into a stable monolithic substance suitable for landfill disposal.

Unlike washed and clean aggregates used in cement concrete, contaminated soil and sludge waste including river sediments can be very heterogeneous with a mixture of materials of different particle size distributions, shapes, densities and surface properties (eg clayey and plastic). Such dissimilar characteristics and rheological properties can compromise the immobilisation treatment.

The responsible person must use a properly designed and engineered treatment plant that is adequately equipped with automatic or semi automatic control in respect of waste and chemical reagents handling. Avoid or prohibit any manual operation which is prone to human error and may be unreliable. A typical process flow diagram for the treatment of contaminated soil and sludge waste is in <u>Annex A</u>.

The mechanical mixer functions as the chemical reactor of the immobilisation treatment process. It should be equipped with a stationary mixing compartment and an agitator fitted with heavy duty mixing paddles/blades; and it can perform the following mixing duties:

- Provide positive agitation/stirring of the mix and achieve rigorous mixing e.g. turbulent flow within the mixing compartment.
- Adequately handle homogeneous and heterogeneous solids including soil, aggregates and sludges, and materials exhibiting plastic properties e.g. silt and clay.
- · Capable of achieving a homogenous mix within minutes of mixing.

As a matter of DECC policy the tumbler type mixer e.g. small DIY rotating concrete mixer, rotating mixer mounted on a delivery truck, rotary hoe or bull dozer are not acceptable mixing devices for the immobilisation treatment of contaminated soil and sludge waste. Such machineries cannot discharge the above mixing duties for processing hazardous waste or sludge.

The following types of mixer (Perry, Chemical Engineers Handbook, McGraw Hill) could attain the above mixing duties and are considered suitable for the purposes of the immobilisation treatment of contaminated soil or sludge waste. They are commonly used in industry operations.

- 1. Pug mill mixer.
- 2. Paddle type mixer including Ribbon mixer and Turbine/Pan mixer.

However, the choice of mixer is a waste specific issue and the responsible person should conduct a test run before adopting the equipment for full scale treatment. The DECC would consider and approve other types of mechanical mixing device on merits.

Annex A: Typical Process Flow Diagram for Contaminated Soil Treatment Plant

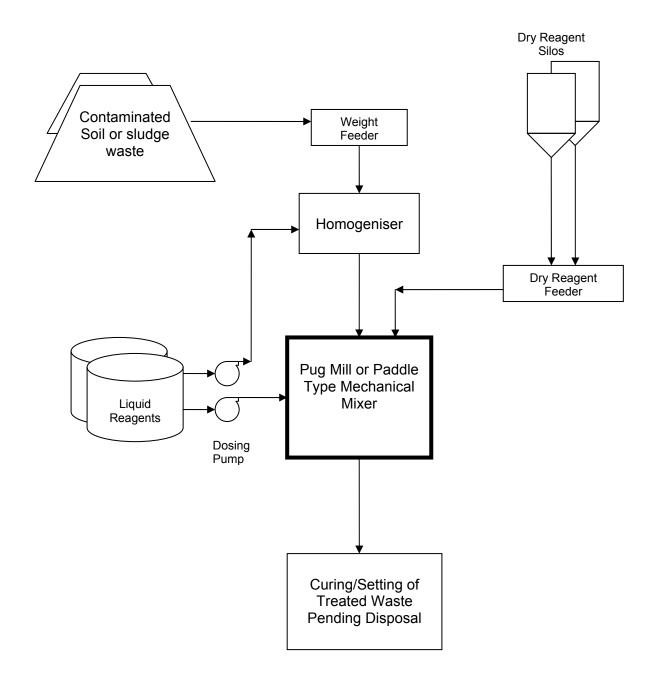
Page last updated: 26 February 2008

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DECC WASTE TECHINICAL NOTE 1: ANNEX A

Typical Process Flow Diagram for Contaminated Soil Treatment Plant





Appendix F

Laboratory Certificates of Analysis



Envirolab Services Pty Ltd ABN 37 112 535 645 12 Ashley St Chatswood NSW 2067 ph 02 9910 6200 fax 02 9910 6201 enquiries@envirolabservices.com.au www.envirolabservices.com.au

CERTIFICATE OF ANALYSIS 42976

<u>Client:</u> JBS Environmental P.O. Box 940 MASCOT NSW 1460

Attention: Tim Davis / Sumi Dorairoj

Sample log in details:

Your Reference: No. of samples: Date samples received: Date completed instructions received:

40913, Macdonaldtown

21 Soils, 1 Water 02/07/10 02/07/10

Analysis Details:

Please refer to the following pages for results, methodology summary and quality control data. Samples were analysed as received from the client. Results relate specifically to the samples as received. Results are reported on a dry weight basis for solids and on an as received basis for other matrices. *Please refer to the last page of this report for any comments relating to the results.*

Report Details: Date results requested by: 9/07/10 Date of Preliminary Report: Not issued Issue Date: 9/07/10 NATA accreditation number 2901. This document shall not be reproduced except in full. This document is issued in accordance with NATA's accreditation requirements. Accredited for compliance with ISO/IEC 17025. Tests not covered by NATA are denoted with *.

Results Approved By:

David Springer/

Business Development & Quality Manager

Envirolab Reference:

Revision No:

42976

R 00

₹a, Sandra Taxlor Senior Organic Chemist

Kluign Morgen

Rhian Morgan Metals Supervisor

Nancy Zhang Chemist

M. Mauffell

Matt Mansfield Approved Signatory

Page 1 of 51

VOCs in soil						
Our Reference:	UNITS	42976-1	42976-2	42976-5	42976-6	42976-7
Your Reference		JBS TP1	JBS TP2	JBS TP2	JBS TP3	JBS TP4
Depth		0.3-0.4	0.4-0.5	1.4-1.5	1.7	0.5
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	06/07/2010	06/07/2010	06/07/2010	06/07/2010	06/07/2010
Date analysed	-	07/07/2010	07/07/2010	07/07/2010	07/07/2010	07/07/2010
Dichlorodifluoromethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Chloromethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Vinyl Chloride	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Bromomethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Chloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Trichlorofluoromethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1-Dichloroethene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
trans-1,2-dichloroethene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1-dichloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
cis-1,2-dichloroethene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
bromochloromethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
chloroform	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,2-dichloropropane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,1-trichloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1-dichloropropene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Cyclohexane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
carbon tetrachloride	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Benzene	mg/kg	<0.5	<0.5	1.4	0.9	<0.5
dibromomethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichloropropane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
trichloroethene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
bromodichloromethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
trans-1,3-dichloropropene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
cis-1,3-dichloropropene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,2-trichloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Toluene	mg/kg	<0.5	<0.5	2.4	0.72	<0.5
1,3-dichloropropane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
dibromochloromethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dibromoethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
tetrachloroethene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,1,2-tetrachloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
chlorobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Ethylbenzene	mg/kg	<1.0	<1.0	26	22	<1.0
bromoform	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
m+p-xylene	mg/kg	<2.0	<2.0	44	13	<2.0
styrene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,2,2-tetrachloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0

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VOCs in soil						
Our Reference:	UNITS	42976-1	42976-2	42976-5	42976-6	42976-7
Your Reference		JBS TP1	JBS TP2	JBS TP2	JBS TP3	JBS TP4
Depth		0.3-0.4	0.4-0.5	1.4-1.5	1.7	0.5
Date Sampled Type of sample		2/07/2010 Soil	2/07/2010 Soil	2/07/2010 Soil	2/07/2010 Soil	2/07/2010 Soil
o-Xylene	mg/kg	<1.0	<1.0	22	18	<1.0
1,2,3-trichloropropane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
isopropylbenzene	mg/kg	<1.0	<1.0	4.4	3.3	<1.0
bromobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
n-propyl benzene	mg/kg	<1.0	<1.0	5.7	3.2	<1.0
2-chlorotoluene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
4-chlorotoluene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,3,5-trimethyl benzene	mg/kg	<1.0	<1.0	25	15	<1.0
tert-butyl benzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,4-trimethyl benzene	mg/kg	<1.0	<1.0	54	36	<1.0
1,3-dichlorobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
sec-butyl benzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,4-dichlorobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
4-isopropyl toluene	mg/kg	<1.0	<1.0	2.7	2.5	<1.0
1,2-dichlorobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
n-butyl benzene	mg/kg	<1.0	<1.0	3.2	1.9	<1.0
1,2-dibromo-3-chloropropane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,4-trichlorobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
hexachlorobutadiene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,3-trichlorobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Surrogate Dibromofluorometha	%	110	108	106	110	106
Surrogate aaa-Trifluorotoluene	%	110	112	90	109	122
Surrogate Toluene-d	%	89	83	69	86	86
Surrogate 4-Bromofluorobenzene	%	100	100	118	105	101

VOCs in soil						
Our Reference:	UNITS	42976-8	42976-9	42976-14	42976-16	42976-17
Your Reference		JBS TP4	JBS TP4	JBS TP5	JBS TP5	JBS TP5
Depth		1.0	1.6-1.7	0.5	1.5	2.0
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	06/07/2010	06/07/2010	06/07/2010	06/07/2010	06/07/2010
Date analysed	-	07/07/2010	07/07/2010	07/07/2010	07/07/2010	07/07/2010
Dichlorodifluoromethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Chloromethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Vinyl Chloride	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Bromomethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Chloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Trichlorofluoromethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1-Dichloroethene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
trans-1,2-dichloroethene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1-dichloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
cis-1,2-dichloroethene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
bromochloromethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
chloroform	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,2-dichloropropane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,1-trichloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1-dichloropropene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Cyclohexane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
carbon tetrachloride	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Benzene	mg/kg	<0.5	<0.5	<0.5	<0.5	<0.5
dibromomethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichloropropane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
trichloroethene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
bromodichloromethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
trans-1,3-dichloropropene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
cis-1,3-dichloropropene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,2-trichloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Toluene	mg/kg	<0.5	<0.5	<0.5	<0.5	<0.5
1,3-dichloropropane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
dibromochloromethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dibromoethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
tetrachloroethene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,1,2-tetrachloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
chlorobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Ethylbenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
bromoform	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
m+p-xylene	mg/kg	<2.0	<2.0	<2.0	5.3	<2.0
styrene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,2,2-tetrachloroethane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1, 1, 2, 2-15 li a 5 li 01 06 li la 116	iiig/Kg	>1.0	×1.0	NI.0	NI.U	×1.0

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VOCs in soil						
Our Reference:	UNITS	42976-8	42976-9	42976-14	42976-16	42976-17
Your Reference		JBS TP4	JBS TP4	JBS TP5	JBS TP5	JBS TP5
Depth		1.0	1.6-1.7	0.5	1.5	2.0
Date Sampled Type of sample		2/07/2010 Soil	2/07/2010 Soil	2/07/2010 Soil	2/07/2010 Soil	2/07/2010 Soil
o-Xylene	mg/kg	<1.0	<1.0	<1.0	3.3	<1.0
1,2,3-trichloropropane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
isopropylbenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
bromobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
n-propyl benzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2-chlorotoluene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
4-chlorotoluene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,3,5-trimethyl benzene	mg/kg	<1.0	<1.0	<1.0	4.1	<1.0
tert-butyl benzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,4-trimethyl benzene	mg/kg	<1.0	<1.0	<1.0	8.5	1.7
1,3-dichlorobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
sec-butyl benzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,4-dichlorobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
4-isopropyl toluene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichlorobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
n-butyl benzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dibromo-3-chloropropane	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,4-trichlorobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
hexachlorobutadiene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,3-trichlorobenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Surrogate Dibromofluorometha	%	112	113	104	112	114
Surrogate aaa-Trifluorotoluene	%	104	92	87	90	108
Surrogate Toluene-da	%	76	66	66	66	78
Surrogate 4-Bromofluorobenzene	%	100	100	100	101	100

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VOCs in soil Our Reference:	UNITS	42976-19	42976-21
Your Reference		QA2	JBS TP3
Depth		-	4.0-4.2
Date Sampled		2/07/2010	2/07/2010
Type of sample		Soil	Soil
Date extracted	-	06/07/2010	06/07/2010
Date analysed	-	07/07/2010	07/07/2010
Dichlorodifluoromethane	mg/kg	<1.0	<1.0
Chloromethane	mg/kg	<1.0	<1.0
Vinyl Chloride	mg/kg	<1.0	<1.0
Bromomethane	mg/kg	<1.0	<1.0
Chloroethane	mg/kg	<1.0	<1.0
Trichlorofluoromethane	mg/kg	<1.0	<1.0
1,1-Dichloroethene	mg/kg	<1.0	<1.0
trans-1,2-dichloroethene	mg/kg	<1.0	<1.0
1,1-dichloroethane	mg/kg	<1.0	<1.0
cis-1,2-dichloroethene	mg/kg	<1.0	<1.0
bromochloromethane	mg/kg	<1.0	<1.0
chloroform	mg/kg	<1.0	<1.0
2,2-dichloropropane	mg/kg	<1.0	<1.0
1,2-dichloroethane	mg/kg	<1.0	<1.0
1,1,1-trichloroethane	mg/kg	<1.0	<1.0
1,1-dichloropropene	mg/kg	<1.0	<1.0
Cyclohexane	mg/kg	<1.0	<1.0
carbon tetrachloride	mg/kg	<1.0	<1.0
Benzene	mg/kg	<0.5	<0.5
dibromomethane	mg/kg	<1.0	<1.0
1,2-dichloropropane	mg/kg	<1.0	<1.0
trichloroethene	mg/kg	<1.0	<1.0
bromodichloromethane	mg/kg	<1.0	<1.0
trans-1,3-dichloropropene	mg/kg	<1.0	<1.0
cis-1,3-dichloropropene	mg/kg	<1.0	<1.0
1,1,2-trichloroethane	mg/kg	<1.0	<1.0
Toluene	mg/kg	<0.5	<0.5
1,3-dichloropropane	mg/kg	<1.0	<1.0
dibromochloromethane	mg/kg	<1.0	<1.0
1,2-dibromoethane	mg/kg	<1.0	<1.0
tetrachloroethene	mg/kg	<1.0	<1.0
1,1,1,2-tetrachloroethane	mg/kg	<1.0	<1.0
chlorobenzene	mg/kg	<1.0	<1.0
Ethylbenzene	mg/kg	<1.0	<1.0
bromoform	mg/kg	<1.0	<1.0
m+p-xylene	mg/kg	<2.0	<2.0
styrene	mg/kg	<1.0	<1.0
1,1,2,2-tetrachloroethane	mg/kg	<1.0	<1.0

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VOCs in soil			
Our Reference:	UNITS	42976-19	42976-21
Your Reference		QA2	JBS TP3
Depth		-	4.0-4.2
Date Sampled		2/07/2010	2/07/2010
Type of sample		Soil	Soil
o-Xylene	mg/kg	<1.0	<1.0
1,2,3-trichloropropane	mg/kg	<1.0	<1.0
isopropylbenzene	mg/kg	<1.0	<1.0
bromobenzene	mg/kg	<1.0	<1.0
n-propyl benzene	mg/kg	<1.0	<1.0
2-chlorotoluene	mg/kg	<1.0	<1.0
4-chlorotoluene	mg/kg	<1.0	<1.0
1,3,5-trimethyl benzene	mg/kg	<1.0	<1.0
tert-butyl benzene	mg/kg	<1.0	<1.0
1,2,4-trimethyl benzene	mg/kg	<1.0	2.2
1,3-dichlorobenzene	mg/kg	<1.0	<1.0
sec-butyl benzene	mg/kg	<1.0	<1.0
1,4-dichlorobenzene	mg/kg	<1.0	<1.0
4-isopropyl toluene	mg/kg	<1.0	<1.0
1,2-dichlorobenzene	mg/kg	<1.0	<1.0
n-butyl benzene	mg/kg	<1.0	<1.0
1,2-dibromo-3-chloropropane	mg/kg	<1.0	<1.0
1,2,4-trichlorobenzene	mg/kg	<1.0	<1.0
hexachlorobutadiene	mg/kg	<1.0	<1.0
1,2,3-trichlorobenzene	mg/kg	<1.0	<1.0
Surrogate Dibromofluorometha	%	116	110
Surrogate aaa-Trifluorotoluene	%	111	89
Surrogate Toluene-da	%	86	66
Surrogate 4-Bromofluorobenzene	%	101	100

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BTEX in Soil			
		10070 11	10070 10
Our Reference:	UNITS	42976-11	42976-12
Your Reference		Trip Spike	Trip Blank
Depth		-	-
Date Sampled		2/07/2010	2/07/2010
Type of sample		Soil	Soil
Date extracted	-	06/07/2010	06/07/2010
Date analysed	-	07/07/2010	07/07/2010
Benzene	mg/kg	98	<0.5
Toluene	mg/kg	100	<0.5
Ethylbenzene	mg/kg	96	<1.0
m+p-xylene	mg/kg	96	<2.0
o-Xylene	mg/kg	100	<1.0
Surrogate aaa-Trifluorotoluene	%	132	137

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PAHs in Soil						
Our Reference:	UNITS	42976-1	42976-2	42976-5	42976-6	42976-7
Your Reference		JBS TP1	JBS TP2	JBS TP2	JBS TP3	JBS TP4
Depth		0.3-0.4	0.4-0.5	1.4-1.5	1.7	0.5
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	06/07/2010	06/07/2010	06/07/2010	06/07/2010	06/07/2010
Date analysed	-	07/07/2010	07/07/2010	07/07/2010	07/07/2010	07/07/2010
Naphthalene	mg/kg	0.6	13	350	310	0.3
Acenaphthylene	mg/kg	0.7	25	6.0	1.3	0.8
Acenaphthene	mg/kg	0.1	3.8	3.7	3.1	<0.1
Fluorene	mg/kg	0.5	22	9.9	2.7	0.2
Phenanthrene	mg/kg	3.4	110	23	4.6	1.4
Anthracene	mg/kg	0.8	32	6.7	1.3	0.5
Fluoranthene	mg/kg	4.4	110	14	1.8	3.1
Pyrene	mg/kg	4.9	130	19	2.3	5.1
Benzo(a)anthracene	mg/kg	2.6	75	7.5	0.7	2.8
Chrysene	mg/kg	2.6	63	6.2	0.7	3.1
Benzo(b+k)fluoranthene	mg/kg	3.8	79	8.0	0.7	5.6
Benzo(a)pyrene	mg/kg	3.0	64	7.7	0.6	4.4
Indeno(1,2,3-c,d)pyrene	mg/kg	1.3	20	2.6	0.2	2.5
Dibenzo(a,h)anthracene	mg/kg	0.3	6.4	0.7	<0.1	0.6
Benzo(g,h,i)perylene	mg/kg	1.1	17	2.7	0.2	2.8
Surrogate p-Terphenyl-d14	%	97	100	96	97	99

PAHs in Soil						
Our Reference:	UNITS	42976-8	42976-9	42976-14	42976-16	42976-17
Your Reference		JBS TP4	JBS TP4	JBS TP5	JBS TP5	JBS TP5
Depth		1.0	1.6-1.7	0.5	1.5	2.0
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	06/07/2010	06/07/2010	06/07/2010	06/07/2010	06/07/2010
Date analysed	-	07/07/2010	07/07/2010	07/07/2010	07/07/2010	07/07/2010
Naphthalene	mg/kg	<0.1	<0.1	6.3	250	9.2
Acenaphthylene	mg/kg	<0.1	<0.1	9.7	22	0.9
Acenaphthene	mg/kg	<0.1	<0.1	2.0	6.4	0.3
Fluorene	mg/kg	<0.1	<0.1	10	31	0.6
Phenanthrene	mg/kg	<0.1	<0.1	39	94	1.7
Anthracene	mg/kg	<0.1	<0.1	11	29	0.5
Fluoranthene	mg/kg	<0.1	<0.1	54	64	1.9
Pyrene	mg/kg	<0.1	<0.1	65	73	2.8
Benzo(a)anthracene	mg/kg	<0.1	<0.1	36	37	1.4
Chrysene	mg/kg	<0.1	<0.1	30	32	1.3
Benzo(b+k)fluoranthene	mg/kg	<0.2	<0.2	36	35	1.8
Benzo(a)pyrene	mg/kg	<0.05	<0.05	30	30	1.5
Indeno(1,2,3-c,d)pyrene	mg/kg	<0.1	<0.1	9.4	9.4	0.5
Dibenzo(a,h)anthracene	mg/kg	<0.1	<0.1	3.0	2.9	0.2
Benzo(g,h,i)perylene	mg/kg	<0.1	<0.1	8.4	8.6	0.5
Surrogate p-Terphenyl-d14	%	93	92	103	91	94

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PAHs in Soil			
Our Reference:	UNITS	42976-19	42976-21
Your Reference		QA2	JBS TP3
Depth		-	4.0-4.2
Date Sampled		2/07/2010	2/07/2010
Type of sample		Soil	Soil
Date extracted	-	06/07/2010	06/07/2010
Date analysed	-	07/07/2010	07/07/2010
Naphthalene	mg/kg	2.8	4.5
Acenaphthylene	mg/kg	1.7	0.6
Acenaphthene	mg/kg	0.2	0.4
Fluorene	mg/kg	0.7	0.8
Phenanthrene	mg/kg	7.9	4.1
Anthracene	mg/kg	2.1	1.0
Fluoranthene	mg/kg	15	4.4
Pyrene	mg/kg	20	5.1
Benzo(a)anthracene	mg/kg	10	2.2
Chrysene	mg/kg	9.1	2.1
Benzo(b+k)fluoranthene	mg/kg	14	2.3
Benzo(a)pyrene	mg/kg	11	1.7
Indeno(1,2,3-c,d)pyrene	mg/kg	4.6	0.6
Dibenzo(a,h)anthracene	mg/kg	1.2	0.2
Benzo(g,h,i)perylene	mg/kg	4.4	0.6
Surrogate p-Terphenyl-d14	%	95	98

Speciated Phenols in Soil						
Our Reference:	UNITS	42976-1	42976-2	42976-5	42976-6	42976-7
Your Reference		JBS TP1	JBS TP2	JBS TP2	JBS TP3	JBS TP4
Depth		0.3-0.4	0.4-0.5	1.4-1.5	1.7	0.5
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	06/07/2010	06/07/2010	06/07/2010	06/07/2010	06/07/2010
Date analysed	-	08/07/2010	08/07/2010	08/07/2010	08/07/2010	08/07/2010
Phenol	mg/kg	<1.0	<10	<1.0	<1.0	<1.0
2-Chlorophenol	mg/kg	<1.0	<10	<1.0	<1.0	<1.0
2-Methylphenol	mg/kg	<1.0	<10	<1.0	<1.0	<1.0
3/4-Methylphenol	mg/kg	<2.0	<20	<2.0	<2.0	<2.0
2-Nitrophenol	mg/kg	<1.0	<10	<1.0	<1.0	<1.0
2,4-Dimethylphenol	mg/kg	<1.0	<10	<1.0	<1.0	<1.0
2,4-Dichlorophenol	mg/kg	<1.0	<10	<1.0	<1.0	<1.0
2,6-dichlorophenol	mg/kg	<1.0	<10	<1.0	<1.0	<1.0
2,4,5-trichlorophenol	mg/kg	<1.0	<10	<1.0	<1.0	<1.0
2,4,6-trichlorophenol	mg/kg	<1.0	<10	<1.0	<1.0	<1.0
2,4-dinitrophenol	mg/kg	<10	<100	<10	<10	<10
4-nitrophenol	mg/kg	<10	<100	<10	<10	<10
2,3,4,6-tetrachlorophenol	mg/kg	<1.0	<10	<1.0	<1.0	<1.0
2-methyl-4,6-dinitrophenol	mg/kg	<10	<100	<10	<10	<10
pentachlorophenol	mg/kg	<10	<100	<10	<10	<10
Surrogate 2-fluorophenol	%	59	68	57	69	81
Surrogate Phenol-d6	%	65	55	57	89	79
Surrogate 2,4,6-Tribromophenol	%	52	85	107	74	66
Surrogate p-Terphenyl-d14	%	103	119	99	103	103

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Speciated Phenols in Soil						
Our Reference:	UNITS	42976-8	42976-9	42976-14	42976-16	42976-17
Your Reference		JBS TP4	JBS TP4	JBS TP5	JBS TP5	JBS TP5
Depth		1.0	1.6-1.7	0.5	1.5	2.0
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	06/07/2010	06/07/2010	06/07/2010	06/07/2010	06/07/2010
Date analysed	-	08/07/2010	08/07/2010	08/07/2010	08/07/2010	08/07/2010
Phenol	mg/kg	<1.0	<1.0	<10	<10	<1.0
2-Chlorophenol	mg/kg	<1.0	<1.0	<10	<10	<1.0
2-Methylphenol	mg/kg	<1.0	<1.0	<10	<10	<1.0
3/4-Methylphenol	mg/kg	<2.0	<2.0	<20	<20	<2.0
2-Nitrophenol	mg/kg	<1.0	<1.0	<10	<10	<1.0
2,4-Dimethylphenol	mg/kg	<1.0	<1.0	<10	<10	<1.0
2,4-Dichlorophenol	mg/kg	<1.0	<1.0	<10	<10	<1.0
2,6-dichlorophenol	mg/kg	<1.0	<1.0	<10	<10	<1.0
2,4,5-trichlorophenol	mg/kg	<1.0	<1.0	<10	<10	<1.0
2,4,6-trichlorophenol	mg/kg	<1.0	<1.0	<10	<10	<1.0
2,4-dinitrophenol	mg/kg	<10	<10	<100	<100	<10
4-nitrophenol	mg/kg	<10	<10	<100	<100	<10
2,3,4,6-tetrachlorophenol	mg/kg	<1.0	<1.0	<10	<10	<1.0
2-methyl-4,6-dinitrophenol	mg/kg	<10	<10	<100	<100	<10
pentachlorophenol	mg/kg	<10	<10	<100	<100	<10
Surrogate 2-fluorophenol	%	88	79	78	76	61
Surrogate Phenol-de	%	77	84	80	88	77
Surrogate 2,4,6-Tribromophenol	%	55	39	103	99	60
Surrogate p-Terphenyl-d14	%	98	101	108	111	101

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40913, Macdonaldtown

Speciated Phenols in Soil			
Our Reference:	UNITS	42976-19	42976-21
Your Reference		QA2	JBS TP3
Depth		-	4.0-4.2
Date Sampled		2/07/2010	2/07/2010
Type of sample		Soil	Soil
Date extracted	-	06/07/2010	06/07/2010
Date analysed	-	08/07/2010	08/07/2010
Phenol	mg/kg	<1.0	<1.0
2-Chlorophenol	mg/kg	<1.0	<1.0
2-Methylphenol	mg/kg	<1.0	<1.0
3/4-Methylphenol	mg/kg	<2.0	<2.0
2-Nitrophenol	mg/kg	<1.0	<1.0
2,4-Dimethylphenol	mg/kg	<1.0	<1.0
2,4-Dichlorophenol	mg/kg	<1.0	<1.0
2,6-dichlorophenol	mg/kg	<1.0	<1.0
2,4,5-trichlorophenol	mg/kg	<1.0	<1.0
2,4,6-trichlorophenol	mg/kg	<1.0	<1.0
2,4-dinitrophenol	mg/kg	<10	<10
4-nitrophenol	mg/kg	<10	<10
2,3,4,6-tetrachlorophenol	mg/kg	<1.0	<1.0
2-methyl-4,6-dinitrophenol	mg/kg	<10	<10
pentachlorophenol	mg/kg	<10	<10
Surrogate 2-fluorophenol	%	76	74
Surrogate Phenol-de	%	87	73
Surrogate 2,4,6-Tribromophenol	%	68	69
Surrogate p-Terphenyl-d14	%	103	106

Client Reference:	40913, Macdonaldtown
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Acid Extractable metals in soil						
Our Reference:	UNITS	42976-1	42976-2	42976-5	42976-6	42976-7
Your Reference		JBS TP1	JBS TP2	JBS TP2	JBS TP3	JBS TP4
Depth		0.3-0.4	0.4-0.5	1.4-1.5	1.7	0.5
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date digested	-	07/07/2010	07/07/2010	07/07/2010	07/07/2010	07/07/2010
Date analysed	-	07/07/2010	07/07/2010	07/07/2010	07/07/2010	07/07/2010
Arsenic	mg/kg	30	13	<4	<4	8
Cadmium	mg/kg	1.1	<0.5	<0.5	<0.5	<0.5
Chromium	mg/kg	26	19	14	12	22
Copper	mg/kg	230	80	4	1	46
Lead	mg/kg	220	220	58	14	260
Mercury	mg/kg	0.3	0.4	0.1	<0.1	0.2
Nickel	mg/kg	20	26	5	2	10
Zinc	mg/kg	260	220	200	3	24
Acid Extractable metals in soil						
Our Reference:	UNITS	42976-8	42976-9	42976-14	42976-16	42976-17
Your Reference		JBS TP4	JBS TP4	JBS TP5	JBS TP5	JBS TP5
Depth		1.0	1.6-1.7	0.5	1.5	2.0
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date digested	-	07/07/2010	07/07/2010	07/07/2010	07/07/2010	07/07/2010
Date analysed	-	07/07/2010	07/07/2010	07/07/2010	07/07/2010	07/07/201
Arsenic	mg/kg	9	6	6	5	6
Cadmium	mg/kg	<0.5	<0.5	<0.5	<0.5	<0.5
Chromium	mg/kg	35	7	18	23	25
Copper	mg/kg	51	65	18	16	9
Lead	mg/kg	61	100	58	50	36
Mercury	mg/kg	<0.1	<0.1	0.1	0.3	<0.1
Nickel	mg/kg	3	4	5	13	7
Zinc	mg/kg	14	47	35	93	27

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Acid Extractable metals in soil			
Our Reference:	UNITS	42976-19	42976-21
Your Reference		QA2	JBS TP3
Depth		-	4.0-4.2
Date Sampled		2/07/2010	2/07/2010
Type of sample		Soil	Soil
Date digested	-	07/07/2010	07/07/2010
Date analysed	-	07/07/2010	07/07/2010
Arsenic	mg/kg	30	5
Cadmium	mg/kg	1.1	<0.5
Chromium	mg/kg	21	22
Copper	mg/kg	260	6
Lead	mg/kg	280	24
Mercury	mg/kg	0.4	<0.1
Nickel	mg/kg	24	3
Zinc	mg/kg	330	9



Moisture						
Our Reference:	UNITS	42976-1	42976-2	42976-5	42976-6	42976-7
Your Reference		JBS TP1	JBS TP2	JBS TP2	JBS TP3	JBS TP4
Depth		0.3-0.4	0.4-0.5	1.4-1.5	1.7	0.5
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date prepared	-	6/7/2010	6/7/2010	6/7/2010	6/7/2010	6/7/2010
Date analysed	-	6/7/2010	6/7/2010	6/7/2010	6/7/2010	6/7/2010
Moisture	%	13	10	25	23	16
Moisture						
Our Reference:	UNITS	42976-8	42976-9	42976-14	42976-16	42976-17
Your Reference		JBS TP4	JBS TP4	JBS TP5	JBS TP5	JBS TP5
Depth		1.0	1.6-1.7	0.5	1.5	2.0
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date prepared	-	6/7/2010	6/7/2010	6/7/2010	6/7/2010	6/7/2010
Date analysed	-	6/7/2010	6/7/2010	6/7/2010	6/7/2010	6/7/2010
Moisture	%	23	17	19	23	27
Moisture				7		
Our Reference:	UNITS	42976-19	42976-21			
Your Reference		QA2	JBS TP3			
Depth		-	4.0-4.2			
Date Sampled		2/07/2010	2/07/2010			
Turna of comple		Call	Cell			

Soil

6/7/2010

6/7/2010

17

-

-

%

Soil

6/7/2010

6/7/2010

25

Type of sample

Date prepared

Date analysed

Moisture

Asbestos ID - soils						
Our Reference:	UNITS	42976-1	42976-2	42976-5	42976-6	42976-7
Your Reference		JBS TP1	JBS TP2	JBS TP2	JBS TP3	JBS TP4
Depth		0.3-0.4	0.4-0.5	1.4-1.5	1.7	0.5
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date analysed	-	9/7/2010	9/7/2010	9/7/2010	9/7/2010	9/7/2010
Sample Description	-	Approx 40g Soil				
Asbestos ID in soil	-	No asbestos found at reporting limit of 0.1g/kg				
Trace Analysis	-	Respirable fibres not detected				
Asbestos ID - soils						
Our Reference:	UNITS	42976-8	42976-9	42976-16	42976-17	42976-21
Your Reference		JBS TP4	JBS TP4	JBS TP5	JBS TP5	JBS TP3
Depth		1.0	1.6-1.7	1.5	2.0	4.0-4.2
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date analysed	-	9/7/2010	9/7/2010	9/7/2010	9/7/2010	9/7/2010
Sample Description	-	Approx 40g Soil				
Asbestos ID in soil	-	No asbestos found at reporting limit of 0.1g/kg				
Trace Analysis	-	Respirable fibres not detected				

VOCs in Zero Headspace Our Reference:	UNITS	42976-1	42976-2	42976-5	42976-6	42976-7
Your Reference		42976-1 JBS TP1	42976-2 JBS TP2	42976-5 JBS TP2	42976-6 JBS TP3	42976-7 JBS TP4
Depth		0.3-0.4	0.4-0.5	1.4-1.5	1.7	0.5
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	8/7/2010	8/7/2010	8/7/2010	8/7/2010	8/7/2010
Date analysed	-	8/7/2010	8/7/2010	8/7/2010	8/7/2010	8/7/2010
Dichlorodifluoromethane	µg/L	<10	<10	<100	<100	<10
Chloromethane	µg/L	<10	<10	<100	<100	<10
Vinyl Chloride	µg/L	<10	<10	<100	<100	<10
Bromomethane	µg/L	<10	<10	<100	<100	<10
Chloroethane	µg/L	<10	<10	<100	<100	<10
Trichlorofluoromethane	µg/L	<10	<10	<100	<100	<10
1,1-Dichloroethene	µg/L	<1.0	<1.0	<100	<100	<1.0
Trans-1,2-dichloroethene	µg/L	<1.0	<1.0	<100	<100	<1.0
1,1-dichloroethane	µg/L	<1.0	<1.0	<10	<10	<1.0
Cis-1,2-dichloroethene	µg/L	<1.0	<1.0	<10	<10	<1.0
Bromochloromethane	µg/L	<1.0	<1.0	<10	<10	<1.0
Chloroform	µg/L	<1.0	<1.0	<10	<10	<1.0
2,2-dichloropropane	µg/L	<1.0	<1.0	<10	<10	<1.0
1,2-dichloroethane	µg/L	<1.0	<1.0	<10	<10	<1.0
1,1,1-trichloroethane	µg/L	<1.0	<1.0	<10	<10	<1.0
1,1-dichloropropene	µg/L	<1.0	<1.0	<10	<10	<1.0
Carbon tetrachloride	µg/L	<1.0	<1.0	<10	<10	<1.0
Benzene	µg/L	<1.0	<1.0	72	32	<1.0
Dibromomethane	µg/L	<1.0	<1.0	<10	<10	<1.0
1,2-dichloropropane	µg/L	<1.0	<1.0	<10	<10	<1.0
Trichloroethene	µg/L	<1.0	<1.0	<10	<10	<1.0
Bromodichloromethane	µg/L	<1.0	<1.0	<10	<10	<1.0
trans-1,3-dichloropropene	µg/L	<1.0	<1.0	<10	<10	<1.0
cis-1,3-dichloropropene	µg/L	<1.0	<1.0	<10	<10	<1.0
1,1,2-trichloroethane	µg/L	<1.0	<1.0	<10	<10	<1.0
Toluene	µg/L	<1.0	1.7	110	20	<1.0
1,3-dichloropropane	µg/L	<1.0	<1.0	<10	<10	<1.0
Dibromochloromethane	µg/L	<1.0	<1.0	<10	<10	<1.0
1,2-dibromoethane	µg/L	<1.0	<1.0	<10	<10	<1.0
Tetrachloroethene	µg/L	<1.0	<1.0	<10	<10	<1.0
1,1,1,2-tetrachloroethane	µg/L	<1.0	<1.0	<10	<10	<1.0
Chlorobenzene	µg/L	<1.0	<1.0	<10	<10	<1.0
Ethylbenzene	µg/L	<1.0	2.3	630	490	<1.0
Bromoform	µg/L	<1.0	<1.0	<10	<10	<1.0
m+p-xylene	µg/L	<2.0	5.2	980	300	<2.0
Styrene	µg/L	<1.0	<1.0	<10	<10	<1.0
1,1,2,2-tetrachloroethane	µg/L	<1.0	<1.0	<10	<10	<1.0
o-xylene	µg/L	<1.0	2.1	510	430	<1.0

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VOCs in Zero Headspace						
Our Reference:	UNITS	42976-1	42976-2	42976-5	42976-6	42976-7
Your Reference		JBS TP1	JBS TP2	JBS TP2	JBS TP3	JBS TP4
Depth		0.3-0.4	0.4-0.5	1.4-1.5	1.7	0.5
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
1,2,3-trichloropropane	μg/L	<1.0	<1.0	<10	<10	<1.0
Isopropylbenzene	μg/L	<1.0	<1.0	72	58	<1.0
Bromobenzene	μg/L	<1.0	<1.0	<10	<10	<1.0
n-propyl benzene	μg/L	<1.0	<1.0	73	51	<1.0
2-chlorotoluene	μg/L	<1.0	<1.0	<10	<10	<1.0
4-chlorotoluene	μg/L	<1.0	<1.0	<10	<10	<1.0
1,3,5-trimethyl benzene	μg/L	<1.0	1.5	340	260	<1.0
Tert-butyl benzene	μg/L	<1.0	<1.0	<10	<10	<1.0
1,2,4-trimethyl benzene	μg/L	<1.0	2.7	820	640	<1.0
1,3-dichlorobenzene	μg/L	<1.0	<1.0	<10	<10	<1.0
Sec-butyl benzene	μg/L	<1.0	<1.0	<10	<10	<1.0
1,4-dichlorobenzene	μg/L	<1.0	<1.0	<10	<10	<1.0
4-isopropyl toluene	μg/L	<1.0	<1.0	21	28	1.7
1,2-dichlorobenzene	μg/L	<1.0	<1.0	<10	<10	<1.0
n-butyl benzene	μg/L	<1.0	<1.0	14	16	<1.0
1,2-dibromo-3-chloropropane	μg/L	<1.0	<1.0	<10	<10	<1.0
1,2,4-trichlorobenzene	μg/L	<1.0	<1.0	<10	<10	<1.0
Hexachlorobutadiene	µg/L	<1.0	<1.0	<10	<10	<1.0
1,2,3-trichlorobenzene	µg/L	<1.0	<1.0	<10	<10	<1.0
Surrogate Dibromofluoromethane	%	115	#	117	118	#
Surrogate toluene-d8	%	86	125	93	88	137
Surrogate 4-BFB	%	104	106	105	104	115

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Client Reference:

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VOCs in Zero Headspace Our Reference:	UNITS	42976-8	42976-9	42976-14	42976-16	42976-17
Your Reference		JBS TP4	JBS TP4	JBS TP5	JBS TP5	JBS TP5
Depth		1.0	1.6-1.7	0.5	1.5	2.0
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	8/7/2010	8/7/2010	8/7/2010	8/7/2010	8/7/2010
Date analysed	-	8/7/2010	8/7/2010	8/7/2010	8/7/2010	8/7/2010
Dichlorodifluoromethane	µg/L	<10	<10	<10	<10	<10
Chloromethane	µg/L	<10	<10	<10	<10	<10
Vinyl Chloride	µg/L	<10	<10	<10	<10	<10
Bromomethane	µg/L	<10	<10	<10	<10	<10
Chloroethane	µg/L	<10	<10	<10	<10	<10
Trichlorofluoromethane	µg/L	<10	<10	<10	<10	<10
1,1-Dichloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Trans-1,2-dichloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1-dichloroethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Cis-1,2-dichloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Bromochloromethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Chloroform	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
2,2-dichloropropane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichloroethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,1-trichloroethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1-dichloropropene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Carbon tetrachloride	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Benzene	µg/L	<1.0	<1.0	<1.0	<1.0	3.7
Dibromomethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dichloropropane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Trichloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Bromodichloromethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
trans-1,3-dichloropropene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
cis-1,3-dichloropropene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,2-trichloroethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Toluene	µg/L	<1.0	<1.0	<1.0	35	4.8
1,3-dichloropropane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Dibromochloromethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2-dibromoethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Tetrachloroethene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,1,1,2-tetrachloroethane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Chlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Ethylbenzene	μg/L	<1.0	1.2	2.0	17	12
Bromoform	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
m+p-xylene	μg/L	<2.0	2.1	<2.0	350	21
Styrene	μg/L	<1.0	<1.0	<1.0	2.0	<1.0
1,1,2,2-tetrachloroethane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
o-xylene	μg/L	<1.0	<1.0	1.6	220	19
	-	1	1	1	1	

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VOCs in Zero Headspace						
Our Reference:	UNITS	42976-8	42976-9	42976-14	42976-16	42976-17
Your Reference		JBS TP4	JBS TP4	JBS TP5	JBS TP5	JBS TP5
Depth		1.0	1.6-1.7	0.5	1.5	2.0
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
1,2,3-trichloropropane	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Isopropylbenzene	μg/L	<1.0	<1.0	<1.0	<1.0	1.1
Bromobenzene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
n-propyl benzene	μg/L	<1.0	<1.0	<1.0	1.2	1.5
2-chlorotoluene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
4-chlorotoluene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,3,5-trimethyl benzene	μg/L	<1.0	<1.0	1.9	170	11
Tert-butyl benzene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,4-trimethyl benzene	μg/L	<1.0	2.8	5.2	380	31
1,3-dichlorobenzene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Sec-butyl benzene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,4-dichlorobenzene	μg/L	<1.0	<1.0	<1.0	<1.0	<1.0
4-isopropyl toluene	µg/L	<1.0	<1.0	<1.0	15	1.1
1,2-dichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
n-butyl benzene	µg/L	<1.0	<1.0	<1.0	2.1	<1.0
1,2-dibromo-3-chloropropane	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,4-trichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Hexachlorobutadiene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
1,2,3-trichlorobenzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Surrogate Dibromofluoromethane	%	#	#	#	#	111
Surrogate toluene-d8	%	111	94	116	112	89
Surrogate 4-BFB	%	105	113	112	122	108

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VOCs in Zero Headspace			
Our Reference:	UNITS	42976-19	42976-21
Your Reference		QA2	JBS TP3
Depth		-	4.0-4.2
Date Sampled		2/07/2010	2/07/2010
Type of sample		Soil	Soil
Date extracted	-	8/7/2010	8/7/2010
Date analysed	-	8/7/2010	8/7/2010
Dichlorodifluoromethane	µg/L	<10	<10
Chloromethane	µg/L	<10	<10
Vinyl Chloride	µg/L	<10	<10
Bromomethane	µg/L	<10	<10
Chloroethane	µg/L	<10	<10
Trichlorofluoromethane	µg/L	<10	<10
1,1-Dichloroethene	µg/L	<1.0	<1.0
Trans-1,2-dichloroethene	µg/L	<1.0	<1.0
1,1-dichloroethane	µg/L	<1.0	<1.0
Cis-1,2-dichloroethene	µg/L	<1.0	<1.0
Bromochloromethane	µg/L	<1.0	<1.0
Chloroform	μg/L	<1.0	<1.0
2,2-dichloropropane	μg/L	<1.0	<1.0
1,2-dichloroethane	μg/L	<1.0	<1.0
1,1,1-trichloroethane	μg/L	<1.0	<1.0
1,1-dichloropropene	μg/L	<1.0	<1.0
Carbon tetrachloride	μg/L	<1.0	<1.0
Benzene	μg/L	<1.0	1.0
Dibromomethane	µg/L	<1.0	<1.0
1,2-dichloropropane	µg/L	<1.0	<1.0
Trichloroethene	μg/L	<1.0	<1.0
Bromodichloromethane	µg/L	<1.0	<1.0
trans-1,3-dichloropropene	µg/L	<1.0	<1.0
cis-1,3-dichloropropene	µg/L	<1.0	<1.0
1,1,2-trichloroethane	µg/L	<1.0	<1.0
Toluene	µg/L	<1.0	<1.0
1,3-dichloropropane	µg/L	<1.0	<1.0
Dibromochloromethane	µg/L	<1.0	<1.0
1,2-dibromoethane	µg/L	<1.0	<1.0
Tetrachloroethene	µg/L	<1.0	<1.0
1,1,1,2-tetrachloroethane	µg/L	<1.0	<1.0
Chlorobenzene	µg/L	<1.0	<1.0
Ethylbenzene	µg/L	1.7	17
Bromoform	µg/L	<1.0	<1.0
m+p-xylene	µg/L	2.7	7.5
Styrene	µg/L	<1.0	<1.0
1,1,2,2-tetrachloroethane	µg/L	<1.0	<1.0
o-xylene	µg/L	1.6	11



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VOCs in Zero Headspace			
Our Reference:	UNITS	42976-19	42976-21
Your Reference		QA2	JBS TP3
Depth		-	4.0-4.2
Date Sampled		2/07/2010	2/07/2010
Type of sample		Soil	Soil
1,2,3-trichloropropane	µg/L	<1.0	<1.0
Isopropylbenzene	µg/L	<1.0	2.8
Bromobenzene	µg/L	<1.0	<1.0
n-propyl benzene	µg/L	<1.0	3.3
2-chlorotoluene	µg/L	<1.0	<1.0
4-chlorotoluene	µg/L	<1.0	<1.0
1,3,5-trimethyl benzene	µg/L	1.4	29
Tert-butyl benzene	µg/L	<1.0	<1.0
1,2,4-trimethyl benzene	µg/L	3.6	57
1,3-dichlorobenzene	µg/L	<1.0	<1.0
Sec-butyl benzene	µg/L	<1.0	<1.0
1,4-dichlorobenzene	µg/L	<1.0	<1.0
4-isopropyl toluene	µg/L	<1.0	3.4
1,2-dichlorobenzene	µg/L	<1.0	<1.0
n-butyl benzene	µg/L	<1.0	3.3
1,2-dibromo-3-chloropropane	µg/L	<1.0	<1.0
1,2,4-trichlorobenzene	µg/L	<1.0	<1.0
Hexachlorobutadiene	µg/L	<1.0	<1.0
1,2,3-trichlorobenzene	µg/L	<1.0	<1.0
Surrogate Dibromofluoromethane	%	120	114
Surrogate toluene-d8	%	91	90
Surrogate 4-BFB	%	106	111

Client Reference:	40913, Macdonaldtown
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Metals in TCLP USEPA1311						
Our Reference:	UNITS	42976-1	42976-2	42976-5	42976-6	42976-7
Your Reference		JBS TP1	JBS TP2	JBS TP2	JBS TP3	JBS TP
Depth		0.3-0.4	0.4-0.5	1.4-1.5	1.7	0.5
Date Sampled Type of sample		2/07/2010 Soil	2/07/2010 Soil	2/07/2010 Soil	2/07/2010 Soil	2/07/20 ² Soil
Date extracted	-	07/07/2010	07/07/2010	07/07/2010	07/07/2010	07/07/20
Date analysed	-	08/07/2010	08/07/2010	08/07/2010	08/07/2010	08/07/20
pH of soil for fluid# determ.	pH units	7.40	7.50	7.40	7.50	6.70
pH of soil for fluid # determ. (acid)	pH units	1.40	1.40	1.40	1.60	1.40
Extraction fluid used	-	1	1	1	1	1
pH of final Leachate	pH units	5.30	5.00	5.00	5.10	5.00
Arsenic in TCLP	mg/L	<0.05	<0.05	<0.05	<0.05	<0.05
Cadmium in TCLP	mg/L	<0.01	<0.01	<0.01	<0.01	<0.01
Chromium in TCLP	mg/L	<0.01	<0.01	<0.01	<0.01	<0.01
Copper in TCLP	mg/L	0.1	0.04	0.04	0.02	0.1
Lead in TCLP	mg/L	<0.03	0.09	<0.03	<0.03	0.05
Mercury in TCLP	mg/L	<0.0005	<0.0005	<0.0005	<0.0005	<0.000
Nickel in TCLP	mg/L	0.02	<0.02	<0.02	<0.02	<0.02
Zinc in TCLP	mg/L	1.6	1.7	0.5	1.0	1.4
Metals in TCLP USEPA1311		42076.9	42076.0	42076 14	42076 16	42076 1
Our Reference: Your Reference Depth	UNITS	42976-8 JBS TP4 1.0	42976-9 JBS TP4 1.6-1.7	42976-14 JBS TP5 0.5	42976-16 JBS TP5 1.5	JBS TP 2.0
Our Reference: Your Reference	UNITS	JBS TP4	JBS TP4	JBS TP5	JBS TP5	JBS TP 2.0
Our Reference: Your Reference Depth Date Sampled	UNITS 	JBS TP4 1.0 2/07/2010	JBS TP4 1.6-1.7 2/07/2010	JBS TP5 0.5 2/07/2010	JBS TP5 1.5 2/07/2010	JBS TP 2.0 2/07/20 Soil
Our Reference: Your Reference Depth Date Sampled Type of sample	UNITS 	JBS TP4 1.0 2/07/2010 Soil	JBS TP4 1.6-1.7 2/07/2010 Soil	JBS TP5 0.5 2/07/2010 Soil	JBS TP5 1.5 2/07/2010 Soil	JBS TP 2.0 2/07/20 Soil 07/07/20
Our Reference: Your Reference Depth Date Sampled Type of sample Date extracted	UNITS 	JBS TP4 1.0 2/07/2010 Soil 07/07/2010	JBS TP4 1.6-1.7 2/07/2010 Soil 07/07/2010	JBS TP5 0.5 2/07/2010 Soil 07/07/2010	JBS TP5 1.5 2/07/2010 Soil 07/07/2010	JBS TP 2.0 2/07/20 Soil 07/07/20
Our Reference: Your Reference Depth Date Sampled Type of sample Date extracted Date analysed pH of soil for fluid# determ.	 - pH units	JBS TP4 1.0 2/07/2010 Soil 07/07/2010 08/07/2010 5.90	JBS TP4 1.6-1.7 2/07/2010 Soil 07/07/2010 08/07/2010	JBS TP5 0.5 2/07/2010 Soil 07/07/2010 08/07/2010	JBS TP5 1.5 2/07/2010 Soil 07/07/2010 08/07/2010	JBS TP 2.0 2/07/20 Soil 07/07/20 08/07/20 7.00
Our Reference: Your Reference Depth Date Sampled Type of sample Date extracted Date analysed	 - -	JBS TP4 1.0 2/07/2010 Soil 07/07/2010 08/07/2010	JBS TP4 1.6-1.7 2/07/2010 Soil 07/07/2010 08/07/2010 6.10	JBS TP5 0.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.00	JBS TP5 1.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.60	JBS TP 2.0 2/07/20 Soil 07/07/20 08/07/20
Our Reference: Your Reference Depth Date Sampled Type of sample Date extracted Date analysed pH of soil for fluid# determ. pH of soil for fluid # determ. (acid) Extraction fluid used	 - pH units pH units -	JBS TP4 1.0 2/07/2010 Soil 07/07/2010 08/07/2010 5.90 1.40 1	JBS TP4 1.6-1.7 2/07/2010 Soil 07/07/2010 08/07/2010 6.10 1.40 1	JBS TP5 0.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.00 1.30 1	JBS TP5 1.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.60 1.30 1	JBS TP 2.0 2/07/20 Soil 07/07/20 08/07/20 7.00 1.40 1
Our Reference: Your Reference Depth Date Sampled Type of sample Date extracted Date analysed pH of soil for fluid# determ. pH of soil for fluid # determ. (acid) Extraction fluid used pH of final Leachate	 - - pH units pH units - pH units	JBS TP4 1.0 2/07/2010 Soil 07/07/2010 08/07/2010 5.90 1.40 1 4.90	JBS TP4 1.6-1.7 2/07/2010 Soil 07/07/2010 08/07/2010 6.10 1.40 1 5.00	JBS TP5 0.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.00 1.30 1 5.00	JBS TP5 1.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.60 1.30 1 5.10	JBS TP 2.0 2/07/20 Soil 07/07/20 08/07/20 7.00 1.40 1 5.00
Our Reference: Your Reference Depth Date Sampled Type of sample Date extracted Date analysed pH of soil for fluid# determ. pH of soil for fluid # determ. (acid) Extraction fluid used pH of final Leachate Arsenic in TCLP	 - pH units pH units - pH units mg/L	JBS TP4 1.0 2/07/2010 Soil 07/07/2010 08/07/2010 5.90 1.40 1 4.90 <0.05	JBS TP4 1.6-1.7 2/07/2010 Soil 07/07/2010 08/07/2010 6.10 1.40 1 5.00 <0.05	JBS TP5 0.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.00 1.30 1 5.00 <0.05	JBS TP5 1.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.60 1.30 1 5.10 <0.05	JBS TP 2.0 2/07/20 Soil 07/07/20 08/07/20 7.00 1.40 1 5.00 <0.05
Our Reference: Your Reference Depth Date Sampled Type of sample Date extracted Date analysed pH of soil for fluid# determ. pH of soil for fluid # determ. (acid) Extraction fluid used pH of final Leachate Arsenic in TCLP Cadmium in TCLP	 - pH units pH units - pH units mg/L mg/L	JBS TP4 1.0 2/07/2010 Soil 07/07/2010 08/07/2010 5.90 1.40 1 4.90 <0.05 <0.01	JBS TP4 1.6-1.7 2/07/2010 Soil 07/07/2010 08/07/2010 6.10 1.40 1 5.00 <0.05 <0.01	JBS TP5 0.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.00 1.30 1 5.00 <0.05 <0.01	JBS TP5 1.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.60 1.30 1 5.10 <0.05 <0.01	JBS TP 2.0 2/07/20 Soil 07/07/20 08/07/20 7.00 1.40 1 5.00 <0.05 <0.01
Our Reference: Your Reference Depth Date Sampled Type of sample Date extracted Date analysed pH of soil for fluid# determ. pH of soil for fluid # determ. (acid) Extraction fluid used pH of final Leachate Arsenic in TCLP Cadmium in TCLP Chromium in TCLP	 - pH units pH units - pH units mg/L mg/L mg/L	JBS TP4 1.0 2/07/2010 Soil 07/07/2010 08/07/2010 5.90 1.40 1 4.90 <0.05 <0.01 <0.01	JBS TP4 1.6-1.7 2/07/2010 Soil 07/07/2010 08/07/2010 6.10 1.40 1 5.00 <0.05 <0.01 <0.01	JBS TP5 0.5 2/07/2010 Soil 07/07/2010 6.00 1.30 1 5.00 <0.05 <0.01 <0.01	JBS TP5 1.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.60 1.30 1 5.10 <0.05 <0.01 <0.01	JBS TP 2.0 2/07/20 Soil 07/07/20 08/07/20 7.00 1.40 1 5.00 <0.05 <0.01 <0.01
Our Reference: Your Reference Depth Date Sampled Type of sample Date extracted Date analysed pH of soil for fluid# determ. pH of soil for fluid # determ. (acid) Extraction fluid used pH of final Leachate Arsenic in TCLP Cadmium in TCLP Chromium in TCLP	 pH units pH units - pH units mg/L mg/L mg/L mg/L	JBS TP4 1.0 2/07/2010 Soil 07/07/2010 08/07/2010 5.90 1.40 1 4.90 <0.05 <0.01 <0.01 0.06	JBS TP4 1.6-1.7 2/07/2010 Soil 07/07/2010 08/07/2010 6.10 1.40 1 5.00 <0.05 <0.01 <0.01 0.4	JBS TP5 0.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.00 1.30 1 5.00 <0.05 <0.01 <0.01 0.02	JBS TP5 1.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.60 1.30 1 5.10 <0.05 <0.01 <0.01 0.02	JBS TP 2.0 2/07/20 Soil 07/07/20 08/07/20 7.00 1.40 1 5.00 <0.05 <0.01 <0.01 0.02
Our Reference: Your Reference Depth Date Sampled Type of sample Date extracted Date analysed pH of soil for fluid # determ. pH of soil for fluid # determ. (acid) Extraction fluid used pH of final Leachate Arsenic in TCLP Cadmium in TCLP Chromium in TCLP Copper in TCLP Lead in TCLP	 pH units pH units - pH units mg/L mg/L mg/L mg/L mg/L	JBS TP4 1.0 2/07/2010 Soil 07/07/2010 08/07/2010 5.90 1.40 1 4.90 <0.05 <0.01 <0.01 0.06 0.04	JBS TP4 1.6-1.7 2/07/2010 Soil 07/07/2010 08/07/2010 6.10 1.40 1 5.00 <0.05 <0.01 <0.01 0.4 0.06	JBS TP5 0.5 2/07/2010 Soil 07/07/2010 6.00 1.30 1 5.00 <0.05 <0.01 <0.01 0.02 0.05	JBS TP5 1.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.60 1.30 1 5.10 <0.05 <0.01 <0.01 0.02 <0.03	JBS TP 2.0 2/07/20 Soil 07/07/20 08/07/20 7.00 1.40 1 5.00 <0.05 <0.01 <0.01 0.02 0.03
Our Reference: Your Reference Depth Date Sampled Type of sample Date extracted Date analysed pH of soil for fluid# determ. pH of soil for fluid # determ. (acid) Extraction fluid used pH of final Leachate Arsenic in TCLP Cadmium in TCLP Chromium in TCLP	 pH units pH units - pH units mg/L mg/L mg/L mg/L	JBS TP4 1.0 2/07/2010 Soil 07/07/2010 08/07/2010 5.90 1.40 1 4.90 <0.05 <0.01 <0.01 0.06	JBS TP4 1.6-1.7 2/07/2010 Soil 07/07/2010 08/07/2010 6.10 1.40 1 5.00 <0.05 <0.01 <0.01 0.4	JBS TP5 0.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.00 1.30 1 5.00 <0.05 <0.01 <0.01 0.02	JBS TP5 1.5 2/07/2010 Soil 07/07/2010 08/07/2010 6.60 1.30 1 5.10 <0.05 <0.01 <0.01 0.02	2/07/20 Soil 07/07/20 08/07/20 7.00 1.40 1 5.00 <0.05 <0.01 <0.01 0.02



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Metals in TCLP USEPA1311			
Our Reference:	UNITS	42976-19	42976-21
Your Reference		QA2	JBS TP3
Depth		-	4.0-4.2
Date Sampled		2/07/2010	2/07/2010
Type of sample		Soil	Soil
Date extracted	-	07/07/2010	07/07/2010
Date analysed	-	08/07/2010	08/07/2010
pH of soil for fluid# determ.	pH units	6.50	5.90
pH of soil for fluid # determ. (acid)	pH units	1.40	1.40
Extraction fluid used	-	1	1
pH of final Leachate	pH units	5.20	5.00
Arsenic in TCLP	mg/L	<0.05	<0.05
Cadmium in TCLP	mg/L	<0.01	<0.01
Chromium in TCLP	mg/L	<0.01	<0.01
Copper in TCLP	mg/L	0.1	0.02
Lead in TCLP	mg/L	0.07	<0.03
Mercury in TCLP	mg/L	<0.0005	<0.0005
Nickel in TCLP	mg/L	0.05	<0.02
Zinc in TCLP	mg/L	1.8	0.2

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PAHs in TCLP (USEPA 1311)						
Our Reference:	UNITS	42976-1	42976-2	42976-5	42976-6	42976-7
Your Reference		JBS TP1	JBS TP2	JBS TP2	JBS TP3	JBS TP4
Depth		0.3-0.4	0.4-0.5	1.4-1.5	1.7	0.5
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	07/07/2010	07/07/2010	07/07/2010	07/07/2010	07/07/2010
Date analysed	-	08/07/2010	08/07/2010	08/07/2010	08/07/2010	08/07/2010
Naphthalene in TCLP	mg/L	<0.001	0.022	2.8	3.6	0.002
Acenaphthylene in TCLP	mg/L	<0.001	0.002	0.02	0.02	<0.001
Acenaphthene in TCLP	mg/L	<0.001	0.001	0.014	0.047	<0.001
Fluorene in TCLP	mg/L	<0.001	0.004	0.020	0.027	<0.001
Phenanthrene in TCLP	mg/L	<0.001	0.005	0.018	0.020	<0.001
Anthracene in TCLP	mg/L	<0.001	0.001	0.003	0.004	<0.001
Fluoranthene in TCLP	mg/L	<0.001	0.001	0.002	0.002	<0.001
Pyrene in TCLP	mg/L	<0.001	0.001	0.002	0.003	<0.001
Benzo(a)anthracene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Chrysene in TCLP	mg/L	<0.001	<0.001	<0.001	0.001	<0.001
Benzo(b+k)fluoranthene in TCLP	mg/L	<0.002	<0.002	<0.002	<0.002	<0.002
Benzo(a)pyrene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Indeno(1,2,3-c,d)pyrene - TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Dibenzo(a,h)anthracene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Benzo(g,h,i)perylene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Surrogate p-Terphenyl-d14	%	124	119	112	115	122



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PAHs in TCLP (USEPA 1311)						
Our Reference:	UNITS	42976-8	42976-9	42976-14	42976-16	42976-17
Your Reference		JBS TP4	JBS TP4	JBS TP5	JBS TP5	JBS TP5
Depth		1.0	1.6-1.7	0.5	1.5	2.0
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	07/07/2010	07/07/2010	07/07/2010	07/07/2010	07/07/2010
Date analysed	-	08/07/2010	08/07/2010	08/07/2010	08/07/2010	08/07/2010
Naphthalene in TCLP	mg/L	<0.001	<0.001	0.001	0.64	0.18
Acenaphthylene in TCLP	mg/L	<0.001	<0.001	<0.001	0.02	0.003
Acenaphthene in TCLP	mg/L	<0.001	<0.001	<0.001	0.008	0.002
Fluorene in TCLP	mg/L	<0.001	<0.001	0.001	0.020	0.002
Phenanthrene in TCLP	mg/L	<0.001	<0.001	0.001	0.014	0.002
Anthracene in TCLP	mg/L	<0.001	<0.001	<0.001	0.003	<0.001
Fluoranthene in TCLP	mg/L	<0.001	<0.001	<0.001	0.001	0.001
Pyrene in TCLP	mg/L	<0.001	<0.001	<0.001	0.001	0.001
Benzo(a)anthracene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Chrysene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Benzo(b+k)fluoranthene in TCLP	mg/L	<0.002	<0.002	<0.002	<0.002	<0.002
Benzo(a)pyrene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Indeno(1,2,3-c,d)pyrene - TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Dibenzo(a,h)anthracene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Benzo(g,h,i)perylene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Surrogate p-Terphenyl-d14	%	130	76	133	123	129



40913, Macdonaldtown

PAHs in TCLP (USEPA 1311)			
Our Reference:	UNITS	42976-19	42976-21
Your Reference		QA2	JBS TP3
Depth		-	4.0-4.2
Date Sampled		2/07/2010	2/07/2010
Type of sample		Soil	Soil
Date extracted	-	07/07/2010	07/07/2010
Date analysed	-	08/07/2010	08/07/2010
Naphthalene in TCLP	mg/L	0.030	0.066
Acenaphthylene in TCLP	mg/L	<0.001	0.001
Acenaphthene in TCLP	mg/L	0.001	0.004
Fluorene in TCLP	mg/L	0.001	0.006
Phenanthrene in TCLP	mg/L	0.001	0.009
Anthracene in TCLP	mg/L	0.001	0.001
Fluoranthene in TCLP	mg/L	<0.001	0.001
Pyrene in TCLP	mg/L	<0.001	0.001
Benzo(a)anthracene in TCLP	mg/L	<0.001	<0.001
Chrysene in TCLP	mg/L	<0.001	<0.001
Benzo(b+k)fluoranthene in TCLP	mg/L	<0.002	<0.002
Benzo(a)pyrene in TCLP	mg/L	<0.001	<0.001
Indeno(1,2,3-c,d)pyrene - TCLP	mg/L	<0.001	<0.001
Dibenzo(a,h)anthracene in TCLP	mg/L	<0.001	<0.001
Benzo(g,h,i)perylene in TCLP	mg/L	<0.001	<0.001
Surrogate p-Terphenyl-d14	%	117	122

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Phenols in TCLP extract						
Our Reference:	UNITS	42976-1	42976-2	42976-5	42976-6	42976-7
Your Reference		JBS TP1	JBS TP2	JBS TP2	JBS TP3	JBS TP4
Depth		0.3-0.4	0.4-0.5	1.4-1.5	1.7	0.5
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Total Phenolics (as Phenol)	mg/L	<0.050	<0.050	<0.050	<0.050	<0.050

Phenols in TCLP extract						
Our Reference:	UNITS	42976-8	42976-9	42976-14	42976-16	42976-17
Your Reference		JBS TP4	JBS TP4	JBS TP5	JBS TP5	JBS TP5
Depth		1.0	1.6-1.7	0.5	1.5	2.0
Date Sampled		2/07/2010	2/07/2010	2/07/2010	2/07/2010	2/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Total Phenolics (as Phenol)	mg/L	<0.050	<0.050	<0.050	<0.050	<0.050

Phenols in TCLP extract			
Our Reference:	UNITS	42976-19	42976-21
Your Reference		QA2	JBS TP3
Depth		-	4.0-4.2
Date Sampled		2/07/2010	2/07/2010
Type of sample		Soil	Soil
Total Phenolics (as Phenol)	mg/L	<0.050	<0.050



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VOCs in water		
Our Reference:	UNITS	42976-13
Your Reference		Rinsate
Depth		-
Date Sampled Type of sample		2/07/2010 Water
Date extracted	-	6/7/2010
Date analysed	-	6/7/2010
Dichlorodifluoromethane	μg/L	<10
Chloromethane	μg/L	<10
Vinyl Chloride	µg/L	<10
Bromomethane	μg/L	<10
Chloroethane	µg/L	<10
Trichlorofluoromethane	µg/L	<10
1,1-Dichloroethene	μg/L	<1.0
Trans-1,2-dichloroethene	µg/L	<1.0
1,1-dichloroethane	μg/L	<1.0
Cis-1,2-dichloroethene	µg/L	<1.0
Bromochloromethane	µg/L	<1.0
Chloroform	μg/L	<1.0
2,2-dichloropropane	μg/L	<1.0
1,2-dichloroethane	μg/L	<1.0
1,1,1-trichloroethane	μg/L	<1.0
1,1-dichloropropene	μg/L	<1.0
Cyclohexane	μg/L	<1.0
Carbon tetrachloride	µg/L	<1.0
Benzene	µg/L	<1.0
Dibromomethane	µg/L	<1.0
1,2-dichloropropane	µg/L	<1.0
Trichloroethene	µg/L	<1.0
Bromodichloromethane	µg/L	<1.0
trans-1,3-dichloropropene	µg/L	<1.0
cis-1,3-dichloropropene	µg/L	<1.0
1,1,2-trichloroethane	µg/L	<1.0
Toluene	μg/L	<1.0
1,3-dichloropropane	μg/L	<1.0
Dibromochloromethane	μg/L	<1.0
1,2-dibromoethane	µg/L	<1.0
Tetrachloroethene	µg/L	<1.0
1,1,1,2-tetrachloroethane	µg/L	<1.0
Chlorobenzene	μg/L	<1.0
Ethylbenzene	µg/L	<1.0
Bromoform	μg/L	<1.0
m+p-xylene	µg/L	<2.0
Styrene	μg/L	<1.0
1,1,2,2-tetrachloroethane	µg/L	<1.0



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VOCs in water Our Reference: Your Reference Depth	UNITS	42976-13 Rinsate -
Date Sampled Type of sample		2/07/2010 Water
o-xylene	µg/L	<1.0
1,2,3-trichloropropane	μg/L	<1.0
Isopropylbenzene	μg/L	<1.0
Bromobenzene	μg/L	<1.0
n-propyl benzene	µg/L	<1.0
2-chlorotoluene	μg/L	<1.0
4-chlorotoluene	μg/L	<1.0
1,3,5-trimethyl benzene	μg/L	<1.0
Tert-butyl benzene	μg/L	<1.0
1,2,4-trimethyl benzene	μg/L	<1.0
1,3-dichlorobenzene	μg/L	<1.0
Sec-butyl benzene	μg/L	<1.0
1,4-dichlorobenzene	μg/L	<1.0
4-isopropyl toluene	μg/L	<1.0
1,2-dichlorobenzene	μg/L	<1.0
n-butyl benzene	μg/L	<1.0
1,2-dibromo-3-chloropropane	µg/L	<1.0
1,2,4-trichlorobenzene	μg/L	<1.0
Hexachlorobutadiene	μg/L	<1.0
1,2,3-trichlorobenzene	μg/L	<1.0
Surrogate Dibromofluoromethane	%	113
Surrogate toluene-d8	%	98
Surrogate 4-BFB	%	103



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PAHs in Water		
Our Reference:	UNITS	42976-13
Your Reference		Rinsate
Depth		-
Date Sampled		2/07/2010
Type of sample		Water
Date extracted	-	08/07/2010
Date analysed	-	08/07/2010
Naphthalene	µg/L	<1
Acenaphthylene	µg/L	<1
Acenaphthene	µg/L	<1
Fluorene	µg/L	<1
Phenanthrene	µg/L	<1
Anthracene	μg/L	<1
Fluoranthene	µg/L	<1
Pyrene	µg/L	<1
Benzo(a)anthracene	μg/L	<1
Chrysene	µg/L	<1
Benzo(b+k)fluoranthene	µg/L	<2
Benzo(a)pyrene	μg/L	<1
Indeno(1,2,3-c,d)pyrene	μg/L	<1
Dibenzo(a,h)anthracene	μg/L	<1
Benzo(g,h,i)perylene	μg/L	<1
Surrogate p-Terphenyl-d14	%	119

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Speciated Phenols in water		
Our Reference:	UNITS	42976-13
Your Reference		Rinsate
Depth		-
Date Sampled		2/07/2010 Water
Type of sample		water
Date extracted	-	08/07/2010
Date analysed	-	08/07/2010
Phenol	µg/L	<10
2-Chlorophenol	µg/L	<10
2-Methylphenol	μg/L	<10
3/4-Methylphenol	µg/L	<20
2-Nitrophenol	μg/L	<10
2,4-Dimethylphenol	μg/L	<10
2,4-Dichlorophenol	μg/L	<10
2,6-Dichlorophenol	μg/L	<10
2,4,5-Trichlorophenol	μg/L	<10
2,4,6-Trichlorophenol	μg/L	<10
2,4-Dinitrophenol	µg/L	<100
4-Nitrophenol	μg/L	<100
2,3,4,6-Tetrachlorophenol	μg/L	<10
2-methyl-4,6-dinitrophenol	μg/L	<100
Pentachlorophenol	μg/L	<100
Surrogate 2-fluorophenol	%	43
Surrogate Phenol-de	%	31
Surrogate 2,4,6-Tribromophenol	%	95
Surrogate p-Terphenyl-d14	%	126

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Metals in Water - Dissolved		
Our Reference:	UNITS	42976-13
Your Reference		Rinsate
Depth		-
Date Sampled		2/07/2010
Type of sample		Water
Date digested	-	09/07/2010
Date analysed	-	09/07/2010
Arsenic - Dissolved	mg/L	<0.05
Cadmium - Dissolved	mg/L	<0.01
Chromium - Dissolved	mg/L	<0.01
Copper - Dissolved	mg/L	<0.01
Lead - Dissolved	mg/L	<0.03
Mercury - Dissolved	mg/L	<0.0005
Nickel - Dissolved	mg/L	<0.02
Zinc - Dissolved	mg/L	<0.02



Client Reference: 40913, Macdonaldtown

Method ID	Methodology Summary
GC.14	Soil samples are extracted with methanol and spiked into water prior to analysing by purge and trap GC-MS.
GC.16	Soil samples are extracted with methanol and spiked into water prior to analysing by purge and trap GC-MS. Water samples are analysed directly by purge and trap GC-MS.
GC.12 subset	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-MS.
GC.12	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-MS.
Metals.20 ICP-AES	Determination of various metals by ICP-AES.
Metals.21 CV-AAS	Determination of Mercury by Cold Vapour AAS.
LAB.8	Moisture content determined by heating at 105 deg C for a minimum of 4 hours.
AS4964-2004	Asbestos ID - Qualitative identification of asbestos type fibres in bulk samples using Polarised Light Microscopy and Dispersion Staining Techniques.
GC.13	Water samples are analysed directly by purge and trap GC-MS.
LAB.4	Toxicity Characteristic Leaching Procedure (TCLP).
EXTRACT.7	Toxicity Characteristic Leaching Procedure (TCLP).
LAB.1	pH - Measured using pH meter and electrode in accordance with APHA 20th ED, 4500-H+.
Metals.20 ICP-AES	Determination of various metals by ICP-AES.
Metals.21 CV-AAS	Determination of Mercury by Cold Vapour AAS.
GC.12 subset	Leachates are extracted with Dichloromethane and analysed by GC-MS.
LAB.30	Total Phenolics - determined colorimetrically following disitillation.

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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
VOCs in soil						Base II Duplicate II %RPD		
Date extracted	-			06/07/2 010	42976-1	06/07/2010 06/07/2010	LCS-2	06/07/2010
Date analysed	-			07/07/2 010	42976-1	07/07/2010 07/07/2010	LCS-2	07/07/2010
Dichlorodifluoromethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
Chloromethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
Vinyl Chloride	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
Bromomethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
Chloroethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
Trichlorofluoromethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
1,1-Dichloroethene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
trans-1,2-dichloroethene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
1,1-dichloroethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	LCS-2	91%
cis-1,2-dichloroethene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
bromochloromethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
chloroform	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	LCS-2	102%
2,2-dichloropropane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
1,2-dichloroethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	LCS-2	97%
1,1,1-trichloroethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	LCS-2	95%
1,1-dichloropropene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
Cyclohexane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
carbon tetrachloride	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
Benzene	mg/kg	0.5	GC.14	<0.5	42976-1	<0.5 <0.5	[NR]	[NR]
dibromomethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
1,2-dichloropropane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
trichloroethene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	LCS-2	98%
bromodichloromethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	LCS-2	88%
trans-1,3-dichloropropen e	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
cis-1,3-dichloropropene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
1,1,2-trichloroethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
Toluene	mg/kg	0.5	GC.14	<0.5	42976-1	<0.5 <0.5	[NR]	[NR]
1,3-dichloropropane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
dibromochloromethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	LCS-2	65%
1,2-dibromoethane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
tetrachloroethene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	LCS-2	86%
1,1,1,2-tetrachloroethan e	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
chlorobenzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
Ethylbenzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
bromoform	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
m+p-xylene	mg/kg	2	GC.14	<2.0	42976-1	<2.0 <2.0	[NR]	[NR]
styrene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]
1,1,2,2-tetrachloroethan e	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]

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	Client Reference: 40913, Macdonaldtown											
QUALITY CONTROL VOCs in soil	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results Base II Duplicate II %RPD	Spike Sm#	Spike % Recovery				
o-Xylene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
1,2,3-trichloropropane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
isopropylbenzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
bromobenzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
n-propyl benzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
2-chlorotoluene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
4-chlorotoluene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
1,3,5-trimethyl benzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
tert-butyl benzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
1,2,4-trimethyl benzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
1,3-dichlorobenzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
sec-butyl benzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
1,4-dichlorobenzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
4-isopropyl toluene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
1,2-dichlorobenzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
n-butyl benzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
1,2-dibromo-3-chloropro pane	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
1,2,4-trichlorobenzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
hexachlorobutadiene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
1,2,3-trichlorobenzene	mg/kg	1	GC.14	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]				
<i>Surrogate</i> Dibromofluorometha	%		GC.14	109	42976-1	110 104 RPD: 6	LCS-2	101%				
Surrogate aaa-Trifluorotoluene	%		GC.14	102	42976-1	110 119 RPD: 8	LCS-2	117%				
Surrogate Toluene-da	%		GC.14	86	42976-1	89 88 RPD: 1	LCS-2	85%				
Surrogate 4-Bromofluorobenzene	%		GC.14	102	42976-1	100 101 RPD: 1	LCS-2	103%				

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Client Reference: 40913, Macdonaldtown

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
BTEX in Soil						Base II Duplicate II %RPD		
Date extracted	-			06/07/2 010	[NT]	[NT]	LCS-2	06/07/2010
Date analysed	-			06/07/2 010	[NT]	[NT]	LCS-2	06/07/2010
Benzene	mg/kg	0.5	GC.16	<0.5	[NT]	[NT]	LCS-2	105%
Toluene	mg/kg	0.5	GC.16	<0.5	[NT]	[NT]	LCS-2	88%
Ethylbenzene	mg/kg	1	GC.16	<1.0	[NT]	[NT]	LCS-2	95%
m+p-xylene	mg/kg	2	GC.16	<2.0	[NT]	[NT]	LCS-2	99%
o-Xylene	mg/kg	1	GC.16	<1.0	[NT]	[NT]	LCS-2	105%
<i>Surrogate</i> aaa-Trifluorotoluene	%		GC.16	102	[NT]	[NT]	LCS-2	126%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
PAHs in Soil						Base II Duplicate II %RPD		
Date extracted	-			06/07/2 010	[NT]	[NT]	LCS-2	06/07/2010
Date analysed	-			07/07/2 010	[NT]	[NT]	LCS-2	07/07/2010
Naphthalene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-2	92%
Acenaphthylene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Acenaphthene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Fluorene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-2	104%
Phenanthrene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-2	97%
Anthracene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Fluoranthene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-2	91%
Pyrene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-2	96%
Benzo(a)anthracene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Chrysene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-2	107%
Benzo(b+k)fluoranthene	mg/kg	0.2	GC.12 subset	<0.2	[NT]	[NT]	[NR]	[NR]
Benzo(a)pyrene	mg/kg	0.05	GC.12 subset	<0.05	[NT]	[NT]	LCS-2	108%
Indeno(1,2,3-c,d)pyrene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Dibenzo(a,h)anthracene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Benzo(g,h,i)perylene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]

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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery			
PAHs in Soil						Base II Duplicate II %RPD	_				
Surrogate p-Terphenyl-d14	%		GC.12 subset	108	[NT]	[NT]	LCS-2	102%			
				Diank	Durlianto Sm#	Duralizata regulto	Onite Om#				
QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery			
Speciated Phenols in Soil						Base II Duplicate II %RPD					
Date extracted	-			06/07/2 010	42976-1	06/07/2010 06/07/2010	LCS-2	06/07/2010			
Date analysed	_			08/07/2 010	42976-1	08/07/2010 08/07/2010	LCS-2	08/07/2010			
Phenol	mg/kg	1	GC.12	<1.0	42976-1	<1.0 <1.0	LCS-2	93%			
2-Chlorophenol	mg/kg	1	GC.12	<1.0	42976-1	<1.0 <1.0	LCS-2	95%			
2-Methylphenol	mg/kg	1	GC.12	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]			
3/4-Methylphenol	mg/kg	2	GC.12	<2.0	42976-1	<2.0 <2.0	[NR]	[NR]			
2-Nitrophenol	mg/kg	1	GC.12	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]			
2,4-Dimethylphenol	mg/kg	1	GC.12	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]			
2,4-Dichlorophenol	mg/kg	1	GC.12	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]			
2,6-dichlorophenol	mg/kg	1	GC.12	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]			
2,4,5-trichlorophenol	mg/kg	1	GC.12	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]			
2,4,6-trichlorophenol	mg/kg	1	GC.12	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]			
2,4-dinitrophenol	mg/kg	10	GC.12	<10	42976-1	<10 <10	[NR]	[NR]			
4-nitrophenol	mg/kg	10	GC.12	<10	42976-1	<10 <10	LCS-2	34%			
2,3,4,6-tetrachlorophenol	mg/kg	1	GC.12	<1.0	42976-1	<1.0 <1.0	[NR]	[NR]			
2-methyl-4,6-dinitrophen ol	mg/kg	10	GC.12	<10	42976-1	<10 <10	[NR]	[NR]			
pentachlorophenol	mg/kg	10	GC.12	<10	42976-1	<10 <10	[NR]	[NR]			
Surrogate 2-fluorophenol	%		GC.12	37	42976-1	59 59 RPD: 0	LCS-2	83%			
Surrogate Phenol-d6	%		GC.12	63	42976-1	65 70 RPD: 7	LCS-2	91%			
Surrogate 2,4,6-Tribromophenol	%		GC.12	58	42976-1	52 66 RPD: 24	LCS-2	53%			
Surrogate	%		GC.12	92	42976-1	103 105 RPD: 2	LCS-2	95%			

p-Terphenyl-d14



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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Acid Extractable metals in soil						Base II Duplicate II %RPD		-
Date digested	-			07/07/2 010	42976-1	07/07/2010 07/07/2010	LCS-1	07/07/2010
Date analysed	-			07/07/2 010	42976-1	07/07/2010 07/07/2010	LCS-1	07/07/2010
Arsenic	mg/kg	4	Metals.20 ICP-AES	<4	42976-1	30 34 RPD: 12	LCS-1	106%
Cadmium	mg/kg	0.5	Metals.20 ICP-AES	<0.5	42976-1	1.1 1.3 RPD: 17	LCS-1	103%
Chromium	mg/kg	1	Metals.20 ICP-AES	<1	42976-1	26 29 RPD: 11	LCS-1	105%
Copper	mg/kg	1	Metals.20 ICP-AES	<1	42976-1	230 290 RPD: 23	LCS-1	93%
Lead	mg/kg	1	Metals.20 ICP-AES	<1	42976-1	220 260 RPD: 17	LCS-1	112%
Mercury	mg/kg	0.1	Metals.21 CV-AAS	<0.1	42976-1	0.3 0.4 RPD: 29	LCS-1	115%
Nickel	mg/kg	1	Metals.20 ICP-AES	<1	42976-1	20 21 RPD: 5	LCS-1	108%
Zinc	mg/kg	1	Metals.20 ICP-AES	<1	42976-1	260 270 RPD: 4	LCS-1	97%

QUALITY CONTROL Moisture	UNITS	PQL	METHOD	Blank
Date prepared	-			6/7/201
				0
Date analysed	-			6/7/201 0
Moisture	%	0.1	LAB.8	<0.10

QUALITY CONTROL	UNITS	PQL	METHOD	Blank
Asbestos ID - soils				
Date analysed	-			[NT]

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
VOCs in Zero Headspace						Base II Duplicate II %RPD		
Date extracted	-			8/7/201 0	[NT]	[NT]	LCS-W1	8/7/2010
Date analysed	-			8/7/201 0	[NT]	[NT]	LCS-W1	8/7/2010
Dichlorodifluoromethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Chloromethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Vinyl Chloride	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Bromomethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Chloroethane	μg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Trichlorofluoromethane	μg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]

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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery	
VOCs in Zero Headspace						Base II Duplicate II %RPD			
1,1-Dichloroethene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Trans-1,2-dichloroethen e	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
1,1-dichloroethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	107%	
Cis-1,2-dichloroethene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Bromochloromethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Chloroform	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	113%	
2,2-dichloropropane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
1,2-dichloroethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	104%	
1,1,1-trichloroethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	108%	
1,1-dichloropropene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Carbon tetrachloride	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Dibromomethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
1,2-dichloropropane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Trichloroethene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	101%	
Bromodichloromethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	122%	
trans-1,3-dichloropropen e	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
cis-1,3-dichloropropene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
1,1,2-trichloroethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Toluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
1,3-dichloropropane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Dibromochloromethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	109%	
1,2-dibromoethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Tetrachloroethene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	83%	
1,1,1,2-tetrachloroethan e	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Chlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Ethylbenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Bromoform	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
m+p-xylene	µg/L	2	GC.13	<2.0	[NT]	[NT]	[NR]	[NR]	
Styrene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
1,1,2,2-tetrachloroethan e	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
o-xylene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
1,2,3-trichloropropane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Isopropylbenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Bromobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
n-propyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
2-chlorotoluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
4-chlorotoluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
1,3,5-trimethyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	
Tert-butyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]	

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		Cli	ent Referer	nce: 4	0913, Macdor	aldtown		
QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
VOCs in Zero Headspace						Base II Duplicate II %RPD		
1,2,4-trimethyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,3-dichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Sec-butyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,4-dichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
4-isopropyl toluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,2-dichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
n-butyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,2-dibromo-3-chloropro pane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,2,4-trichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Hexachlorobutadiene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,2,3-trichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Surrogate Dibromofluoromethane	%		GC.13	109	[NT]	[NT]	LCS-W1	118%
Surrogate toluene-d8	%		GC.13	86	[NT]	[NT]	LCS-W1	86%
Surrogate 4-BFB	%		GC.13	102	[NT]	[NT]	LCS-W1	100%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Metals in TCLP USEPA1311						Base II Duplicate II %RPD		-
Date extracted	-			07/07/2 010	42976-2	07/07/2010 07/07/2010	LCS-W1	07/07/2010
Date analysed	-			08/07/2 010	42976-2	08/07/2010 08/07/2010	LCS-W1	08/07/2010
Arsenic in TCLP	mg/L	0.05	Metals.20 ICP-AES	<0.05	42976-2	<0.05 <0.05	LCS-W1	103%
Cadmium in TCLP	mg/L	0.01	Metals.20 ICP-AES	<0.01	42976-2	<0.01 <0.01	LCS-W1	99%
Chromium in TCLP	mg/L	0.01	Metals.20 ICP-AES	<0.01	42976-2	<0.01 <0.01	LCS-W1	98%
Copper in TCLP	mg/L	0.01	Metals.20 ICP-AES	<0.01	42976-2	0.04 0.04 RPD: 0	LCS-W1	100%
Lead in TCLP	mg/L	0.03	Metals.20 ICP-AES	<0.03	42976-2	0.09 0.1 RPD: 11	LCS-W1	94%
Mercury in TCLP	mg/L	0.0005	Metals.21 CV-AAS	<0.000 5	42976-2	<0.0005 <0.0005	LCS-W1	114%
Nickel in TCLP	mg/L	0.02	Metals.20 ICP-AES	<0.02	42976-2	<0.02 <0.02	LCS-W1	97%
Zinc in TCLP	mg/L	0.02	Metals.20 ICP-AES	<0.02	42976-2	1.7 1.6 RPD: 6	LCS-W1	99%



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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
PAHs in TCLP (USEPA 1311)						Base II Duplicate II %RPD		, , ,
Date extracted	-			07/07/2 010	[NT]	[NT]	LCS-W1	07/07/2010
Date analysed	-			08/07/2 010	[NT]	[NT]	LCS-W1	08/07/2010
Naphthalene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	92%
Acenaphthylene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Acenaphthene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Fluorene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	106%
Phenanthrene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	101%
Anthracene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Fluoranthene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	95%
Pyrene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	101%
Benzo(a)anthracene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Chrysene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	111%
Benzo(b+k)fluoranthene in TCLP	mg/L	0.002	GC.12 subset	<0.002	[NT]	[NT]	[NR]	[NR]
Benzo(a)pyrene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	112%
Indeno(1,2,3-c,d)pyrene - TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Dibenzo(a,h)anthracene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Benzo(g,h,i)perylene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Surrogate p-Terphenyl-d ₁₄	%		GC.12	120	[NT]	[NT]	LCS-W1	124%



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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Phenols in TCLP extract						Base II Duplicate II %RPD		Recovery
Total Phenolics (as Phenol)	mg/L	0.05	LAB.30	<0.050	42976-1	<0.050 <0.050	LCS-W1	91%
QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
VOCs in water						Base II Duplicate II %RPD		,
Date extracted	-			6/7/201 0	[NT]	[NT]	LCS-W1	6/7/2010
Date analysed	-			6/7/201 0	[NT]	[NT]	LCS-W1	6/7/2010
Dichlorodifluoromethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Chloromethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Vinyl Chloride	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Bromomethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Chloroethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
Trichlorofluoromethane	µg/L	10	GC.13	<10	[NT]	[NT]	[NR]	[NR]
1,1-Dichloroethene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Trans-1,2-dichloroethen e	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,1-dichloroethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	100%
Cis-1,2-dichloroethene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Bromochloromethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Chloroform	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	103%
2,2-dichloropropane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,2-dichloroethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	102%
1,1,1-trichloroethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	94%
1,1-dichloropropene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Cyclohexane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Carbon tetrachloride	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Dibromomethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,2-dichloropropane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Trichloroethene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	98%
Bromodichloromethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	77%
trans-1,3-dichloropropen e	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
cis-1,3-dichloropropene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,1,2-trichloroethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Toluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,3-dichloropropane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Dibromochloromethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	64%
1,2-dibromoethane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Tetrachloroethene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	90%
1,1,1,2-tetrachloroethan e	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]

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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike %
VOCs in water						Base II Duplicate II %RPD		Recovery
						·		
Chlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Ethylbenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Bromoform	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
m+p-xylene	µg/L	2	GC.13	<2.0	[NT]	[NT]	[NR]	[NR]
Styrene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,1,2,2-tetrachloroethan e	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
o-xylene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
1,2,3-trichloropropane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Isopropylbenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Bromobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
n-propyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
2-chlorotoluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
4-chlorotoluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
1,3,5-trimethyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
Tert-butyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
1,2,4-trimethyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
1,3-dichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
Sec-butyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
1,4-dichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
4-isopropyl toluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
1,2-dichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
n-butyl benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
1,2-dibromo-3-chloropro pane	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
1,2,4-trichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
Hexachlorobutadiene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
1,2,3-trichlorobenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR
Surrogate Dibromofluoromethane	%		GC.13	106	[NT]	[NT]	LCS-W1	1089
Surrogate toluene-d8	%		GC.13	87	[NT]	[NT]	LCS-W1	89%
Surrogate 4-BFB	%		GC.13	102	[NT]	[NT]	LCS-W1	96%



Client Reference: 40913, Macdonaldtown

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
PAHs in Water						Base II Duplicate II %RPD		, , , , , , , , , , , , , , , , , , ,
Date extracted	-			08/07/2 010	[NT]	[NT]	LCS-W1	08/07/2010
Date analysed	-			08/07/2 010	[NT]	[NT]	LCS-W1	08/07/2010
Naphthalene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	LCS-W1	92%
Acenaphthylene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	[NR]	[NR]
Acenaphthene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	[NR]	[NR]
Fluorene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	LCS-W1	105%
Phenanthrene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	LCS-W1	85%
Anthracene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	[NR]	[NR]
Fluoranthene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	LCS-W1	90%
Pyrene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	LCS-W1	100%
Benzo(a)anthracene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	[NR]	[NR]
Chrysene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	LCS-W1	107%
Benzo(b+k)fluoranthene	µg/L	2	GC.12 subset	<2	[NT]	[NT]	[NR]	[NR]
Benzo(a)pyrene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	LCS-W1	105%
Indeno(1,2,3-c,d)pyrene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	[NR]	[NR]
Dibenzo(a,h)anthracene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	[NR]	[NR]
Benzo(g,h,i)perylene	µg/L	1	GC.12 subset	<1	[NT]	[NT]	[NR]	[NR]
<i>Surrogate</i> p-Terphenyl-d ₁₄	%		GC.12 subset	131	[NT]	[NT]	LCS-W1	133%



Client Reference: 40913, Macdonaldtown

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Speciated Phenols in water						Base II Duplicate II %RPD		
Date extracted	-			08/07/2 010	[NT]	[NT]	LCS-W1	08/07/2010
Date analysed	-			08/07/2 010	[NT]	[NT]	LCS-W1	08/07/2010
Phenol	µg/L	10	GC.12	<10	[NT]	[NT]	LCS-W1	38%
2-Chlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	LCS-W1	83%
2-Methylphenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
3/4-Methylphenol	µg/L	20	GC.12	<20	[NT]	[NT]	[NR]	[NR]
2-Nitrophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4-Dimethylphenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4-Dichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,6-Dichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4,5-Trichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4,6-Trichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4-Dinitrophenol	µg/L	100	GC.12	<100	[NT]	[NT]	[NR]	[NR]
4-Nitrophenol	µg/L	100	GC.12	<100	[NT]	[NT]	LCS-W1	42%
2,3,4,6-Tetrachlorophen ol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2-methyl-4,6-dinitrophen ol	µg/L	100	GC.12	<100	[NT]	[NT]	[NR]	[NR]
Pentachlorophenol	µg/L	100	GC.12	<100	[NT]	[NT]	[NR]	[NR]
Surrogate 2-fluorophenol	%		GC.12	48	[NT]	[NT]	LCS-W1	50%
Surrogate Phenol-d ₆	%		GC.12	31	[NT]	[NT]	LCS-W1	37%
Surrogate 2,4,6-Tribromophenol	%		GC.12	92	[NT]	[NT]	LCS-W1	103%
Surrogate p-Terphenyl-d14	%		GC.12	116	[NT]	[NT]	LCS-W1	113%



Client Reference: 40913, Macdonaldtown

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Metals in Water - Dissolved						Base II Duplicate II %RPD		
Date digested	-			09/07/2 010	[NT]	[NT]	LCS-W1	09/07/2010
Date analysed	-			09/07/2 010	[NT]	[NT]	LCS-W1	09/07/2010
Arsenic - Dissolved	mg/L	0.05	Metals.20 ICP-AES	<0.05	[NT]	[NT]	LCS-W1	94%
Cadmium - Dissolved	mg/L	0.01	Metals.20 ICP-AES	<0.01	[NT]	[NT]	LCS-W1	97%
Chromium - Dissolved	mg/L	0.01	Metals.20 ICP-AES	<0.01	[NT]	[NT]	LCS-W1	94%
Copper - Dissolved	mg/L	0.01	Metals.20 ICP-AES	<0.01	[NT]	[NT]	LCS-W1	95%
Lead - Dissolved	mg/L	0.03	Metals.20 ICP-AES	<0.03	[NT]	[NT]	LCS-W1	94%
Mercury - Dissolved	mg/L	0.0005	Metals.21 CV-AAS	<0.000 5	[NT]	[NT]	LCS-W1	102%
Nickel - Dissolved	mg/L	0.02	Metals.20 ICP-AES	<0.02	[NT]	[NT]	LCS-W1	96%
Zinc - Dissolved	mg/L	0.02	Metals.20 ICP-AES	<0.02	[NT]	[NT]	LCS-W1	95%
QUALITY CONTROL	UNITS	S I	Dup. Sm#		Duplicate	Spike Sm#	Spike % Recovery	
Speciated Phenols in Soil				Base + I	Duplicate + %RPD			
Date extracted	_		[NT]		[NT]	42976-2	06/07/2010	
Date analysed	_		[NT]		[NT]	42976-2	08/07/2010	
Phenol	mg/kg	a	[NT]		[NT]	42976-2	113%	
2-Chlorophenol	mg/kg	a	[NT]		[NT]	42976-2	99%	
2-Methylphenol	mg/kg		[NT]		[NT]	[NR]	[NR]	
3/4-Methylphenol	mg/kg		[NT]		[NT]	[NR]	[NR]	
2-Nitrophenol	mg/kg		[NT]		[NT]	[NR]	[NR]	
2,4-Dimethylphenol	mg/kg	g	[NT]		[NT]	[NR]	[NR]	
2,4-Dichlorophenol	mg/kg	3	[NT]		[NT]	[NR]	[NR]	
2,6-dichlorophenol	mg/kg	3	[NT]		[NT]	[NR]	[NR]	
2,4,5-trichlorophenol	mg/kg	3	[NT]		[NT]	[NR]	[NR]	
2,4,6-trichlorophenol	mg/kg	3	[NT]		[NT]	[NR]	[NR]	
2,4-dinitrophenol	mg/kg		[NT]		[NT]	[NR]	[NR]	
4-nitrophenol	mg/kg	9	[NT]		[NT]	42976-2	42%	
2,3,4,6-tetrachlorophenol	mg/kg	9	[NT]		[NT]	[NR]	[NR]	
2-methyl-4,6-dinitrophenol	l mg/kg	9	[NT]		[NT]	[NR]	[NR]	
pentachlorophenol	mg/kg	9	[NT]		[NT]	[NR]	[NR]	
Surrogate 2-fluorophenol	%		[NT]		[NT]	42976-2	70%	
Surrogate Phenol-d6	%		[NT]		[NT]	42976-2	88%	
<i>Surrogate</i> 2,4,6-Tribromophenol	%		[NT]		[NT]	42976-2	122%	

Envirolab Reference: 42976 Revision No: R 00 ACCREDITED FOR TECHNICAL COMPETENCE

		Client Reference	ce: 40913, Macdonald	town	
QUALITY CONTROL Speciated Phenols in Soil	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery
Surrogate p-Terphenyl-d ₁₄	%	[NT]	[NT]	42976-2	104%
QUALITY CONTROL Acid Extractable metals in soil	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery
Date digested	-	[NT]	[NT]	42976-2	07/07/2010
Date analysed	-	[NT]	[NT]	42976-2	07/07/2010
Arsenic	mg/kg	[NT]	[NT]	42976-2	95%
Cadmium	mg/kg	[NT]	[NT]	42976-2	95%
Chromium	mg/kg	[NT]	[NT]	42976-2	95%
Copper	mg/kg	[NT]	[NT]	42976-2	83%
Lead	mg/kg	[NT]	[NT]	42976-2	107%
Mercury	mg/kg	[NT]	[NT]	42976-2	111%
Nickel	mg/kg	[NT]	[NT]	42976-2	89%
Zinc	mg/kg	[NT]	[NT]	42976-2	94%
QUALITY CONTROL Phenols in TCLP extract	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery
Total Phenolics (as Phenol)	mg/L	[NT]	[NT]	42976-2	115%
QUALITY CONTROL Metals in Water - Dissolved	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery
Date digested	-	[NT]	[NT]	42976-13	09/07/2010
Date analysed	-	[NT]	[NT]	42976-13	09/07/2010
Arsenic - Dissolved	mg/L	[NT]	[NT]	42976-13	96%
Cadmium - Dissolved	mg/L	[NT]	[NT]	42976-13	99%
Chromium - Dissolved	mg/L	[NT]	[NT]	42976-13	94%
Copper - Dissolved	mg/L	[NT]	[NT]	42976-13	96%
Lead - Dissolved	mg/L	[NT]	[NT]	42976-13	95%
Mercury - Dissolved	mg/L	[NT]	[NT]	42976-13	102%
Nickel - Dissolved	mg/L	[NT]	[NT]	42976-13	97%
Zinc - Dissolved	mg/L	[NT]	[NT]	42976-13	96%



Report Comments:

Phenols in soil:PQL has been raised due to the sample matrix requiring dilution.

Asbestos was analysed by Approved Identifier: Matt Mansfield

Asbestos was authorised by Approved Signatory: Matt Mansfield

INS: Insufficient sample for this testNT: Not testedPQL: Practical Quantitation Limit<: Less than</th>>: Greater thanRPD: Relative Percent DifferenceNA: Test not requiredLCS: Laboratory Control SampleNR: Not requested

Quality Control Definitions

Blank: This is the component of the analytical signal which is not derived from the sample but from reagents, glassware etc, can be determined by processing solvents and reagents in exactly the same manner as for samples. **Duplicate**: This is the complete duplicate analysis of a sample from the process batch. If possible, the sample selected should be one where the analyte concentration is easily measurable.

Matrix Spike: A portion of the sample is spiked with a known concentration of target analyte. The purpose of the matrix spike is to monitor the performance of the analytical method used and to determine whether matrix interferences exist.

LCS (Laboratory Control Sample): This comprises either a standard reference material or a control matrix (such as a blank sand or water) fortified with analytes representative of the analyte class. It is simply a check sample.

Surrogate Spike: Surrogates are known additions to each sample, blank, matrix spike and LCS in a batch, of compounds which are similar to the analyte of interest, however are not expected to be found in real samples.

Laboratory Acceptance Criteria:

Duplicate sample and matrix spike recoveries may not be reported on smaller jobs, however, were analysed at a frequency to meet or exceed NEPM requirements. All samples are tested in batches of 20. The duplicate sample RPD and matrix spike recoveries for the sample batch were within laboratory acceptance criteria.

Duplicates: <5xPQL - any RPD is acceptable; >5xPQL - 0-50% RPD is acceptable.

Matrix Spikes and LCS: Generally 70-130% for inorganics/metals; 60-140% for organics and 10-140% for

SVOC and speciated phenols is acceptable. Surrogates: 60-140% is acceptable for general organics and 10-140% for

Envirolab Reference: 429 Revision No: R 0





Envirolab Services Pty Ltd ABN 37 112 535 645 12 Ashley St Chatswood NSW 2067 ph 02 9910 6200 fax 02 9910 6201 enquiries@envirolabservices.com.au www.envirolabservices.com.au

SAMPLE RECEIPT ADVICE

<u>Client:</u> JBS Environmental P.O. Box 940 MASCOT NSW 1460	ph: 8338 1013 Fax: 8338 1700
Attention: Tim Davis / Sumi Dorairoj	
Sample log in details: Your reference: Envirolab Reference: Date received: Date results expected to be reported:	40913, Macdonaldtown 42976 02/07/10 9/07/10

Samples received in appropriate condition for analysis:	YES
No. of samples provided	21 Soils, 1 Water
Turnaround time requested:	Standard
Temperature on receipt	Cool
Cooling Method:	Ice

Comments:

Samples will be held for 1 month for water samples and 2 months for soil samples from date of receipt of samples.

Contact details:

Please direct any queries to Aileen Hie or Jacinta Hurst ph: 02 9910 6200 fax: 02 9910 6201 email: ahie@envirolabservices.com.au or jhurst@envirolabservices.com.au 4125

CHAIN OF CUSTODY - Client

ENVIROLAB SERVICES



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Client: J	BS Environmental			Client	t Proje	ct Nam		Numbe	er:					Envi	rolab	Ser	vic	es	
Project Mgr: Dorairay Sampler: Tim Davis				40913							12 Ashley St, Chatswood, NSW, 2067								
				PO No.:															
Address:	128 O'Risrdon St	-		Envir	olab Se	ervices	Quote	No. :						Phone	e: 02 9	9958	580	1	
	Mascot NSW 20	20		Date	results	requir	red:							Fax:	02 9	9958	580)3	
Email: Sda	orairo j@jbsgroup.con	n.au #tolavi	verbegroup.com	Or ch	oose	standa	rd 7	day / 2	2 day /	3 day				E-mai	il: tno	otaras	s@e	nviro	labservices.com.au
	v =	، Fax: 8338		Note: 1	'nform la rge appli	b in adva	ance if ui	gent tun	naround	is require	ed -			Conta	ict: Ta	nia N	lota	rac	
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		an Aufrick (Market) I	i an		I	}				5 Mai, Isikuwa na we		ner sen m nut fissisiet i							
Envirolab Sample ID	Client Sample ID	Date sampled	Type of sample	Heavy Metals	PAILS	VOCS	Speciated phenols	CLP Hewy Metals	CLP PAHS	VOLS	CLP Dhenols	د -دم + 8TEX	HOLU	ASBestos					Provide as much information about the sample as you can
	JES TP1 /0:3-0:4 M	02/7/10	Seil	₩ ✓			<i>∽</i>						,						Expect high
	JBS TP2 /0.4-0.5 M		1	\checkmark			\checkmark	~	\checkmark	\checkmark	\checkmark								PAtts in all
	JBS TP2/08-0:9 M						1						/						samples
	JBS TP2/12-1-3m			w									\checkmark						
2	JBS TP2/14-15M			\checkmark	$\overline{\checkmark}$		\checkmark	\checkmark		$\overline{}$									
Ŀ	JBS TP3/1-7m							\checkmark	\checkmark								1		
-7	JBS TP 4/05M			\checkmark								г .		$\overline{}$			FINITA		Invirolab Sandices 12 Ashley St Rewood NSW 2067 Ph: 9940 5200
8	JBS TP4/1.0m			 ✓ 				\checkmark	V	~			W			7		Chai	13wood NSW 2067 Ph: 9910,6200
9	JBS TP4/1.6-1.7M					\checkmark	$\overline{}$		1				•			J		d:	42976
١Ũ	JBS TP4/19-20M									1			\checkmark					- 1. A A	2.7
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ιΖ	Trip Blank															r.	amp:¢	oly Ambl	es nt
13	Rinsate	02/07/10	N	 ✓ 	\checkmark	1	~	W	JUN-	week						2	ooling		
+	JBS TP5/0.5m		3	\checkmark			\checkmark	\checkmark			\checkmark							P	Terminong
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Print Name: Spini Doraira				Print Name: 名								Temperature Recieved at: (if applicable)							
		5-30		Date & Time: 2 7 1830								Transp	orted b	y: Ha	nđ de	elivere	d / courier		
Signature: 3				Signa	ture:	•													Page No: 1.4 2

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CHAIN OF CUSTODY - Client

ENVIROLAB SERVICES



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Client: J	<u>BS</u> Environmental			Client	Proje	t Nam					•						ervice		
Project Mgr: Doraira				40913 Macdonalltown						12 A	shley	St, C	Chatsv	vood	, NSW, 2067				
Sampler:	Tim Davis			PO No.: Envirolab Services Quote No. :						Phone: 02 9958 5801									
ddress:	128 O'Riordon St Mascot NSW 20	00						No. :											
			han in	Date results required: Fax: 02 9958 5803 Q U Or choose: standard 1 day / 2 day / 3 day E-mail: tnotaras@enviro Note: Inform lab in advance if urgent turnaround is required - Fax: 02 9958 5803								lahaaniisaa sama.							
	airaj@jbsgroup.com.au/	TOLOUISE	usgroup.com.au								C-IIId	UI	IOLAI	aswe	IIVIIO	abservices.com.au			
Phone: 83		Fax: 833		surchar	ge applie	25											Nota		
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Envirolab Sample ID	Client Sample ID	Date sampled	Type of sample	Heavy metals	PA+IS	1065	speciated phenols	TCLD metals	TELP PAHS	TLP VOCS	TCLD phrnols	Ce-Cq BTEX	070H	Ashestos					Provide as much information about the sample as you can
16	JBSTPJ/15m	02/07/10	S	\sim	\checkmark	 ✓ 	\checkmark	1	V,	\checkmark									
17	JBS TP5 /2.0m		Ì			\checkmark		~	\checkmark	\checkmark									
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Signature:	UC			Signa	cure:									L					Page No: \checkmark°

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ANALYTICAL REPORT

14 July 2010

JBS Environmental Pty Ltd

PO Box 940 MASCOT NSW 1460

Attention:	Sumi Dorairaj		
Your Reference:	40913 - Macdonaldtown		
Our Reference:	SE79619	Samples: Received:	1 Soil 5/7/10
Preliminary Report S	Sent: 12/07/2010	10001100	0,,,10

These samples were analysed in accordance with your written instructions.

For and on Behalf of: SGS ENVIRONMENTAL SERVICES

Sample Receipt: **Production Manager:** Angela Mamalicos Huong Crawford

AU.SampleReceipt.Sydney@sgs.com Huong.Crawford@sgs.com

Results Approved and/or Authorised by:

Elward ibrahun

Edward Ibrahim Laboratory Manager

mlu/ Ly Kim Ha

Organics Signatory



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SGS Australia Pty Ltd Environmental Services Unit 16/33 Maddox Street Alexandria NSW 2015 Australia t +61 (0)2 8594 0400 f + 61 (0)2 8594 0499 ABN 44 000 964 278

www.au.sgs.com

VOCs in Soil - 72 List		
Our Reference:	UNITS	SE79619-1
Your Reference		QC2
Sample Matrix		Soil
Date Sampled		2/07/2010
Date Extracted		8/07/2010
Date Analysed		8/07/2010
Dichlorodifluoromethane(CFC-12)	mg/kg	<1
Chloromethane	mg/kg	<1
Vinyl Chloride (chloroethene)	mg/kg	<0.1
Bromomethane	mg/kg	<1
Chloroethane	mg/kg	<1
Trichlorofluoromethane	mg/kg	<1
Acetone (2-propanone)	mg/kg	<10
1,1-Dichloroethene	mg/kg	<0.1
Methyl lodine (iodomethane)	mg/kg	<5
Acrylonitrile	mg/kg	<0.1
Methylene Chloride (DCM)	mg/kg	<0.5
Allyl Chloride	mg/kg	<0.1
Carbon Disulphide	mg/kg	<0.5
trans-1,2-Dichloroethene	mg/kg	<0.1
Methyl-tert-butyl ether (MtBE)	mg/kg	<0.5
1,1-Dichloroethane	mg/kg	<0.1
2-Butanone (MEK)	mg/kg	<10
cis-1,2-Dichloroethene	mg/kg	<0.1
Bromochloromethane	mg/kg	<0.1
Chloroform	mg/kg	<0.1
2,2-Dichloropropane	mg/kg	<0.1
1,2-Dichloroethane	mg/kg	<0.1
1,1,1-Trichloroethane	mg/kg	<0.1
1,1-Dichloropropene	mg/kg	<0.1
Carbon tetrachloride	mg/kg	<0.1
Benzene	mg/kg	<0.1
Dibromomethane	mg/kg	<0.1
1,2-Dichloropropane	mg/kg	<0.1
Trichloroethene (TCE)	mg/kg	<0.1
2-Nitropropane	mg/kg	<10
Bromodichloromethane	mg/kg	<0.1
cis-1,3-Dichloropropene	mg/kg	<0.1
4-Methyl-2-Pentanone(MIBK)	mg/kg	<1
trans-1,3-Dichloropropene	mg/kg	<0.1
1,1,2-Trichloroethane	mg/kg	<0.1
Toluene	mg/kg	<0.1



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PROJECT: 40913 - Macdonaldtown

VOCs in Soil - 72 List		0570640.4
Our Reference: Your Reference	UNITS	SE79619-1 QC2
Sample Matrix		Soil
Date Sampled		2/07/2010
1,3-Dichloropropane	mg/kg	<0.1
2-Hexanone (MBK)	mg/kg	<5
Dibromochloromethane	mg/kg	<0.1
1,2-Dibromoethane (EDB)	mg/kg	<0.1
Tetrachloroethene (PCE-perchloroethylen	mg/kg	<0.1
1,1,1,2-Tetrachloroethane	mg/kg	<0.1
Chlorobenzene	mg/kg	<0.1
Ethyl benzene	mg/kg	0.1
Bromoform	mg/kg	<0.1
<i>m/p</i> -Xylenes	mg/kg	0.2
Cis-1,4-dichloro-2-butene	mg/kg	<1
Styrene (vinyl benzene)	mg/kg	<0.1
1,1,2,2-Tetrachloroethane	mg/kg	<0.1
o-Xylene	mg/kg	0.1
1,2,3-Trichloropropane	mg/kg	<0.1
Trans-1,4-dichloro-2-butene	mg/kg	<1
Isopropylbenzene (Cumene)	mg/kg	<0.1
Bromobenzene	mg/kg	<0.1
n-Propylbenzene	mg/kg	<0.1
2-Chlorotoluene	mg/kg	<0.1
4-Chlorotoluene	mg/kg	<0.1
1,3,5-Trimethylbenzene	mg/kg	0.1
<i>tert</i> -Butylbenzene	mg/kg	<0.1
1,2,4-Trimethylbenzene	mg/kg	0.3
sec-Butylbenzene	mg/kg	<0.1
1,3-Dichlorobenzene	mg/kg	<0.1
1,4-Dichlorobenzene	mg/kg	<0.1
p-Isopropyl toluene	mg/kg	<0.1
1,2-Dichlorobenzene	mg/kg	<0.1
n-Butylbenzene	mg/kg	<0.1
1,2-Dibromo-3-chloropropane	mg/kg	<0.1
1,2,4-Trichlorobenzene	mg/kg	<0.1
Naphthalene	mg/kg	5.2
Hexachlorobutadiene	mg/kg	<0.1
1,2,3-Trichlorobenzene	mg/kg	<0.1
Vinyl acetate	mg/kg	<10
Dibromofluoromethane	%	94
1,2-Dichloroethane-d4	%	97
Toluene-d8 Surrogate 2	%	83



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PROJECT: 40913 - Macdonaldtown

VOCs in Soil - 72 List		
Our Reference:	UNITS	SE79619-1
Your Reference		QC2
Sample Matrix		Soil
Date Sampled		2/07/2010
4-Bromofluorobenzene Surrogate 3	%	78



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PAHs in Soil		
Our Reference:	UNITS	SE79619-1
Your Reference		QC2
Sample Matrix		Soil
Date Sampled		2/07/2010
Date Extracted		8/07/2010
Date Analysed		8/07/2010
Naphthalene	mg/kg	3.1
2-Methylnaphthalene	mg/kg	1.5
1-Methylnaphthalene	mg/kg	1.1
Acenaphthylene	mg/kg	2.4
Acenaphthene	mg/kg	0.23
Fluorene	mg/kg	0.92
Phenanthrene	mg/kg	5.7
Anthracene	mg/kg	2.6
Fluoranthene	mg/kg	12
Pyrene	mg/kg	13
Benzo[a]anthracene	mg/kg	11
Chrysene	mg/kg	6.3
Benzo[b,k]fluoranthene	mg/kg	12
Benzo[a]pyrene	mg/kg	6.7
Indeno[123-cd]pyrene	mg/kg	4.3
Dibenzo[ah]anthracene	mg/kg	1.2
Benzo[ghi]perylene	mg/kg	4.6
Total PAHs (sum)	mg/kg	89
Nitrobenzene-d5	%	107
2-Fluorobiphenyl	%	103
p -Terphenyl-d14	%	114



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PROJECT: 40913 - Macdonaldtown

1	
UNITS	SE79619-1
	QC2
	Soil
	2/07/2010
	8/07/2010
	8/07/2010
mg/kg	<0.5
mg/kg	<0.5
mg/kg	<1
mg/kg	<0.5
mg/kg	<1
mg/kg	<0.5
mg/kg	<0.5
%	64
%	61
	mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg mg/kg



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PROJECT: 40913 - Macdonaldtown

Metals in Soil by ICP-OES		
Our Reference:	UNITS	SE79619-1
Your Reference		QC2
Sample Matrix		Soil
Date Sampled		2/07/2010
Date Extracted (Metals)		12/06/2010
Date Analysed (Metals)		12/06/2010
Arsenic	mg/kg	45
Cadmium	mg/kg	2.1
Chromium	mg/kg	23
Copper	mg/kg	380
Lead	mg/kg	290
Nickel	mg/kg	24
Zinc	mg/kg	370



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Mercury Cold Vapor/Hg Analyser		
Our Reference:	UNITS	SE79619-1
Your Reference		QC2
Sample Matrix		Soil
Date Sampled		2/07/2010
Date Extracted (Mercury)		8/07/2010
Date Analysed (Mercury)		8/07/2010
Mercury	mg/kg	0.26



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VOCs by ZHE TCLP - 72 List		
Our Reference:	UNITS	SE79619-1
Your Reference		QC2
Sample Matrix		Soil
Date Sampled		2/07/2010
Date Extracted-ZHE TCLP Prep		8/07/2010
pH of soil for fluid# determ.	pH units	8.03
pH of soil for fluid # determ. (acid)	pH units	1.76
Extraction fluid used	-	1
pH of final Leachate	pH units	5.13
Date extracted (VOCs)		8/07/2010
Date analysed (VOCs)		8/07/2010
Dichlorodifluoromethane(CFC-12)	µg/L	<5
Chloromethane	µg/L	<5
Vinyl Chloride (chloroethene)	µg/L	<0.3
Bromomethane	µg/L	<10
Chloroethane	µg/L	<5
Trichlorofluoromethane	μg/L	<1
Acetone (2-propanone)	μg/L	<10
1,1-Dichloroethene	μg/L	<0.5
Methyl lodine (iodomethane)	μg/L	<5
Acrylonitrile	μg/L	<0.5
Methylene Chloride (DCM)	μg/L	<5
Allyl Chloride	μg/L	<0.5
Carbon Disulphide	μg/L	<0.5
trans-1,2-Dichloroethene	μg/L	<0.5
Methyl-tert-butyl ether (MtBE)	μg/L	<2
1,1-Dichloroethane	μg/L	<0.5
2-Butanone (MEK)	μg/L	<10
cis-1,2-Dichloroethene	μg/L	<0.5
Bromochloromethane	μg/L	<0.5
Chloroform	μg/L	<0.5
2,2-Dichloropropane	μg/L	<0.5
1,2-Dichloroethane	μg/L	<0.5
1,1,1-Trichloroethane	μg/L	<0.5
1,1-Dichloropropene	μg/L	<0.5
Carbon tetrachloride	μg/L	<0.5
Benzene	μg/L	<0.5
Dibromomethane	μg/L	<0.5
1,2-Dichloropropane	μg/L	<0.5
Trichloroethene (TCE)	μg/L	<0.5
2-Nitropropane	μg/L	<100
Bromodichloromethane	μg/L	<0.5



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VOCa by 74E TCL B 72 List		
VOCs by ZHE TCLP - 72 List Our Reference:	UNITS	SE79619-1
Your Reference		QC2
Sample Matrix		Soil
Date Sampled		2/07/2010
cis-1,3-Dichloropropene	µg/L	<0.5
4-Methyl-2-Pentanone(MIBK)	µg/L	<5
trans-1,3-Dichloropropene	µg/L	<0.5
1,1,2-Trichloroethane	µg/L	<0.5
Toluene	µg/L	1.1
1,3-Dichloropropane	µg/L	<0.5
2-Hexanone (MBK)	µg/L	<5
Dibromochloromethane	µg/L	<0.5
1,2-Dibromoethane (EDB)	µg/L	<0.5
Tetrachloroethene (PCE-perchloroethylen	µg/L	<0.5
1,1,1,2-Tetrachloroethane	µg/L	<0.5
Chlorobenzene	µg/L	<0.5
Ethyl benzene	μg/L	4.1
Bromoform	μg/L	<0.5
<i>m/p</i> -Xylenes	µg/L	9
Cis-1,4-dichloro-2-butene	µg/L	<1
Styrene (vinyl benzene)	µg/L	<0.5
1,1,2,2-Tetrachloroethane	μg/L	<0.5
o-Xylene	μg/L	6.1
1,2,3-Trichloropropane	µg/L	<0.5
Trans-1,4-dichloro-2-butene	μg/L	<1
Isopropylbenzene (Cumene)	μg/L	0.7
Bromobenzene	µg/L	<0.5
n-Propylbenzene	μg/L	<0.5
2-Chlorotoluene	µg/L	<0.5
4-Chlorotoluene	μg/L	<0.5
1,3,5-Trimethylbenzene	µg/L	4.3
tert-Butylbenzene	µg/L	<0.5
1,2,4-Trimethylbenzene	μg/L	13
sec-Butylbenzene	μg/L	<0.5
1,3-Dichlorobenzene	μg/L	<0.5
1,4-Dichlorobenzene	μg/L	<0.3
p-Isopropyl toluene	µg/L	<0.5
1,2-Dichlorobenzene	μg/L	<0.5
n-Butylbenzene	μg/L	<0.5
1,2-Dibromo-3-chloropropane	μg/L	<0.5
1,2,4-Trichlorobenzene	μg/L	<0.5
Naphthalene	μg/L	45
Hexachlorobutadiene	μg/L	<1



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VOCs by ZHE TCLP - 72 List		
Our Reference:	UNITS	SE79619-1
Your Reference		QC2
Sample Matrix		Soil
Date Sampled		2/07/2010
1,2,3-Trichlorobenzene	µg/L	<0.5
Vinyl acetate	µg/L	<10
Dibromofluoromethane	%	91
1,2-Dichloroethane-d4	%	88
Toluene-d8 Surrogate 2	%	104
4-Bromofluorobenzene Surrogate 3	%	99



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PAHs in TCLP (USEPA 1311) Our Reference: Your Reference Sample Matrix Date Sampled	UNITS 	SE79619-1 QC2 Soil 2/07/2010
Date Extracted (TCLP Preparation)		8/07/2010
pH of soil for fluid# determ.	pH units	8.03
Extraction fluid used	-	1
pH of soil for fluid # determ. (acid)	pH units	1.76
pH of final Leachate	pH units	5.13
Date Extracted		12/07/2010
Date Analysed		12/07/2010
Naphthalene	µg/L	2.1
2-Methylnaphthalene	µg/L	<0.5
1-Methylnaphthalene	µg/L	<0.5
Acenaphthylene	µg/L	<0.50
Acenaphthene	µg/L	<0.50
Fluorene	µg/L	<0.50
Phenanthrene	µg/L	<0.50
Anthracene	µg/L	<0.50
Fluoranthene	µg/L	<0.50
Pyrene	µg/L	<0.50
Benzo[a]anthracene	µg/L	<0.50
Chrysene	µg/L	<0.50
Benzo[b,k]fluoranthene	µg/L	<1.0
Benzo[a]pyrene	µg/L	<0.50
Indeno[123-cd]pyrene	µg/L	<0.50
Dibenzo[ah]anthracene	µg/L	<0.50
Benzo[ghi]perylene	µg/L	<0.50
Total PAHs (sum)	µg/L	<9
Nitrobenzene-d5	%	118
2-Fluorobiphenyl	%	126
p -Terphenyl-d14	%	130



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Metals in TCLP		
Our Reference:	UNITS	SE79619-1
Your Reference		QC2
Sample Matrix		Soil
Date Sampled		2/07/2010
Date Extracted (Metals)		9/07/2010
Date Analysed (Metals)		9/07/2010
Date Extracted (Mercury)		9/07/2010
Date Analysed (Mercury)		9/07/2010
Arsenic	mg/L	<0.05
Cadmium	mg/L	<0.005
Chromium	mg/L	<0.005
Copper	mg/L	0.06
Mercury	mg/L	<0.0001
Nickel	mg/L	0.018
Lead	mg/L	<0.02
Zinc	mg/L	0.42



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Moisture		
Our Reference:	UNITS	SE79619-1
Your Reference		QC2
Sample Matrix		Soil
Date Sampled		2/07/2010
Date Analysed (moisture)		7/06/2010
Moisture	%	17



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Method ID	Methodology Summary
AN433	Volatile Organic Compounds - Soil samples are extracted with methanol, purged and concentrated by a purge and trap apparatus, and then analysed using GC/MS technique. Water samples undergo the same analysis without the extraction step. Based on USEPA 5030B and 8260B.
SEO-019	Volatile Organic Compounds - Soil samples are extracted with methanol, purged and concentrated by a purge and trap apparatus, and then analysed using GC/MS technique. Water samples undergo the same analysis without the extraction step. Based on USEPA 5030B and 8260B.
SEO-030	Polynuclear Aromatic Hydrocarbons - determined by solvent extraction with dichloromethane / acetone for soils and dichloromethane for waters, followed by instrumentation analysis using GC/MS SIM mode.
AN420	Semi-Volatile Organic Compounds (SVOCs) including OC, OP, PCB, Herbicides, PAH, Phthalates, and Speciated Phenols in soils, sediments and waters are determined by GCMS/ECD/FID technique following appropriate solvent extraction process (Based on USEPA 3500C and 8270D).
USEPA 8270	USEPA 8270 GCMS.
SEM-010	Determination of elements by ICP-OES following appropriate sample preparation / digestion process. Based on USEPA 6010C / APHA 21st Edition, 3120B.
SEM-005	Mercury - determined by Cold-Vapour AAS following appropriate sample preparation or digestion process. Based on APHA 21st Edition, 3112B.
AN006	Toxicity Characteristic Leaching Procedure (TCLP) - Preparation of leachates for assessing the mobility of both organic and inorganic contaminants present in liquid, solid, and multiphase wastes. Based on USEPA 1311. For volatile analytes, Zero-Headspace Extraction Vessel (ZHE) is used. This method also meets the requirements of Australian Standard Leaching Procedure (ASLP) AS 4439.3-1997 Part 3.
AN101	pH - Measured using pH meter and electrode based on APHA 21st Edition, 4500-H+. For water analyses the results reported are indicative only as the sample holding time requirement specified in APHA was not met (APHA requires that the pH of the samples are to be measured within 15 minutes after sampling).
SEO-019/SEP-004	Volatile organic contaminants are leached out of the waste with a selected leaching solution in a Zero Headspace Extractor vessel. The leachate is then analysed by Purge and Trap GC/MS technique.
SEO-030	Polynuclear Aromatic Hydrocarbons - determined by solvent extraction with dichloromethane / acetone for soils and dichloromethane for waters, followed by instrumentation analysis using GC/MS SIM mode. Total PAH is the sum of all positive PAH(s) assuming non-quantifiable PAH(s) (<lor) 16="" 50%="" based="" contribute="" do="" for="" is="" lor="" lors.<="" not="" of="" on="" pah="" sum="" td="" the="" to="" total="" total.=""></lor)>
AN002	Preparation of soils, sediments and sludges undergo analysis by either air drying, compositing, subsampling and 1:5 soil water extraction where required. Moisture content is determined by drying the sample at 105 \pm 5°C.



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REPORT NO: SE79619

QUALITY CONTROL	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate Base + Duplicate +	Spike Sm#	Matrix Spike % Recovery Duplicate + %RPD
						%RPD		Duplicate + /// D
Date Extracted				08/07/2 010	[NT]	[NT]	LCS	08/07/2010
Date Analysed				08/07/2 010	[NT]	[NT]	LCS	08/07/2010
Dichlorodifluoromethane (CFC-12)	mg/kg	1.0	AN433	<1	[NT]	[NT]	[NR]	[NR]
Chloromethane	mg/kg	1.0	AN433	<1	[NT]	[NT]	[NR]	[NR]
Vinyl Chloride (chloroethene)	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Bromomethane	mg/kg	1.0	AN433	<1	[NT]	[NT]	[NR]	[NR]
Chloroethane	mg/kg	1.0	AN433	<1	[NT]	[NT]	[NR]	[NR]
Trichlorofluoromethane	mg/kg	1.0	AN433	<1	[NT]	[NT]	[NR]	[NR]
Acetone (2-propanone)	mg/kg	10	AN433	<10	[NT]	[NT]	[NR]	[NR]
1,1-Dichloroethene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	LCS	107%
Methyl lodine (iodomethane)	mg/kg	5	AN433	<5	[NT]	[NT]	[NR]	[NR]
Acrylonitrile	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Methylene Chloride (DCM)	mg/kg	0.5	AN433	<0.5	[NT]	[NT]	[NR]	[NR]
Allyl Chloride	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Carbon Disulphide	mg/kg	0.5	AN433	<0.5	[NT]	[NT]	[NR]	[NR]
trans-1,2-Dichloroeth ene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Methyl-tert-butyl ether (MtBE)	mg/kg	0.5	AN433	<0.5	[NT]	[NT]	[NR]	[NR]
1,1-Dichloroethane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
2-Butanone (MEK)	mg/kg	10	AN433	<10	[NT]	[NT]	[NR]	[NR]
<i>cis</i> -1,2-Dichloroethen e	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Bromochloromethane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Chloroform	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	LCS	79%
2,2-Dichloropropane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,2-Dichloroethane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	LCS	123%
1,1,1-Trichloroethane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,1-Dichloropropene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Carbon tetrachloride	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Benzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	LCS	128%
Dibromomethane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,2-Dichloropropane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Trichloroethene (TCE)	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	LCS	102%
2-Nitropropane	mg/kg	10	AN433	<10	[NT]	[NT]	[NR]	[NR]
Bromodichloromethane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]



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REPORT NO: SE79619

QUALITY CONTROL	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate	Spike Sm#	Matrix Spike % Recovery
VOCs in Soil - 72 List						Base + Duplicate + %RPD		Duplicate + %RPD
<i>cis</i> -1,3-Dichloroprope ne	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
4-Methyl-2-Pentanone (MIBK)	mg/kg	1.0	AN433	<1	[NT]	[NT]	[NR]	[NR]
<i>tran</i> s-1,3-Dichloropro pene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,1,2-Trichloroethane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Toluene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	LCS	116%
1,3-Dichloropropane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
2-Hexanone (MBK)	mg/kg	5	AN433	<5	[NT]	[NT]	[NR]	[NR]
Dibromochloromethane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,2-Dibromoethane (EDB)	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Tetrachloroethene (PCE- perchloroethylen	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,1,1,2-Tetrachloroethan e	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Chlorobenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	LCS	121%
Ethyl benzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	LCS	100%
Bromoform	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
m/p-Xylenes	mg/kg	0.2	AN433	<0.2	[NT]	[NT]	LCS	94%
Cis-1,4-dichloro-2-buten e	mg/kg	1	AN433	<1	[NT]	[NT]	[NR]	[NR]
Styrene (vinyl benzene)	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,1,2,2-Tetrachloroethan e	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
o-Xylene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	LCS	92%
1,2,3-Trichloropropane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Trans-1,4-dichloro-2-but ene	mg/kg	1.0	AN433	<1	[NT]	[NT]	[NR]	[NR]
Isopropylbenzene (Cumene)	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Bromobenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
n-Propylbenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
2-Chlorotoluene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
4-Chlorotoluene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,3,5-Trimethylbenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
tert-Butylbenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,2,4-Trimethylbenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
sec-Butylbenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,3-Dichlorobenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,4-Dichlorobenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
<i>p</i> -Isopropyl toluene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]



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QUALITY CONTROL	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate	Spike Sm#	Matrix Spike % Recovery
VOCs in Soil - 72 List						Base + Duplicate + %RPD		Duplicate + %RPD
1,2-Dichlorobenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
n-Butylbenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,2-Dibromo-3-chloropro pane	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,2,4-Trichlorobenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Naphthalene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Hexachlorobutadiene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
1,2,3-Trichlorobenzene	mg/kg	0.1	AN433	<0.1	[NT]	[NT]	[NR]	[NR]
Vinyl acetate	mg/kg	10	AN433	<10	[NT]	[NT]	[NR]	[NR]
Dibromofluoromethane	%	0	SEO-019	104	[NT]	[NT]	LCS	115%
1,2-Dichloroethane-d4	%	0	SEO-019	113	[NT]	[NT]	LCS	118%
Toluene- <i>d8 Surrogate</i> 2	%	0	SEO-019	77	[NT]	[NT]	LCS	101%
4-Bromofluorobenzene Surrogate 3	%	0	SEO-019	89	[NT]	[NT]	LCS	73%

QUALITY CONTROL PAHs in Soil	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Matrix Spike % Recovery Duplicate + %RPD
Date Extracted				8/07/20 10	[NT]	[NT]	LCS	8/07/2010
Date Analysed				8/07/20 10	[NT]	[NT]	LCS	8/07/2010
Naphthalene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	LCS	86%
2-Methylnaphthalene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	[NR]	[NR]
1-Methylnaphthalene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	[NR]	[NR]
Acenaphthylene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	LCS	93%
Acenaphthene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	LCS	102%
Fluorene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	[NR]	[NR]
Phenanthrene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	LCS	93%
Anthracene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	LCS	98%
Fluoranthene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	LCS	104%
Pyrene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	LCS	107%
Benzo[a]anthracene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	[NR]	[NR]
Chrysene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	[NR]	[NR]
Benzo[<i>b,k</i>]fluoranthe ne	mg/kg	0.2	SEO-030	<0.20	[NT]	[NT]	[NR]	[NR]
Benzo[a]pyrene	mg/kg	0.05	SEO-030	<0.05	[NT]	[NT]	LCS	95%
Indeno[<i>123-cd</i>]pyren e	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	[NR]	[NR]
Dibenzo[<i>ah</i>]anthrace ne	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	[NR]	[NR]



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QUALITY CONTROL PAHs in Soil	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Matrix Spike % Recovery Duplicate + %RPD
Benzo[ghi]perylene	mg/kg	0.1	SEO-030	<0.10	[NT]	[NT]	[NR]	[NR]
Total PAHs (sum)	mg/kg	1.75	SEO-030	<1.7	[NT]	[NT]	[NR]	[NR]
Nitrobenzene-d5	%	0	SEO-030	109	[NT]	[NT]	LCS	96%
2-Fluorobiphenyl	%	0	SEO-030	96	[NT]	[NT]	LCS	93%
p -Terphenyl-d 14	%	0	SEO-030	114	[NT]	[NT]	LCS	118%

QUALITY CONTROL Speciated Phenols in Soil	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Matrix Spike % Recovery Duplicate + %RPD
Date Extracted (Spec. Phenols)				8/07/20 10	[NT]	[NT]	LCS	8/07/2010
Date Analysed (Spec. Phenols)				8/07/20 10	[NT]	[NT]	LCS	8/07/2010
Phenol	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	LCS	75%
2-Methylphenol (o-cresol)	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	[NR]	[NR]
3/4-Methylphenol (m/p-cresol)	mg/kg	1.0	AN420	<1	[NT]	[NT]	[NR]	[NR]
2-Chlorophenol	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	[NR]	[NR]
2.4-Dimethylphenol	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	[NR]	[NR]
2.6-Dichlorophenol	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	LCS	94%
2.4-Dichlorophenol	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	[NR]	[NR]
2-Nitrophenol	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	[NR]	[NR]
2.4.6-Trichlorophenol	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	LCS	72%
4-Nitrophenol	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	[NR]	[NR]
2.4.5-Trichlorophenol	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	[NR]	[NR]
2.3.4.6-Tetrachlorophen ol	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	[NR]	[NR]
Pentachlorophenol	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	LCS	86%
4-Chloro-3-methylphenol	mg/kg	1	AN420	<1	[NT]	[NT]	[NR]	[NR]
2.4-Dinitrophenol	mg/kg	0.5	AN420	<0.5	[NT]	[NT]	[NR]	[NR]
Acetophenone	mg/kg	0.5	USEPA 8270	<0.5	[NT]	[NT]	[NR]	[NR]
2.4.6-Tribromophenol- (Surrogate)	%	10	AN420	110	[NT]	[NT]	LCS	116%
d5-phenol (Surrogate)	%	10	AN420	110	[NT]	[NT]	LCS	116%



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QUALITY CONTROL	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate	Spike Sm#	Matrix Spike % Recovery
Metals in Soil by ICP-OES						Base + Duplicate + %RPD		Duplicate + %RPD
Date Extracted (Metals)				12/06/2 010	[NT]	[NT]	LCS	12/06/2010
Date Analysed (Metals)				12/06/2 010	[NT]	[NT]	LCS	12/06/2010
Arsenic	mg/kg	3	SEM-010	<3	[NT]	[NT]	LCS	99%
Cadmium	mg/kg	0.3	SEM-010	<0.3	[NT]	[NT]	LCS	102%
Chromium	mg/kg	0.3	SEM-010	<0.3	[NT]	[NT]	LCS	103%
Copper	mg/kg	0.5	SEM-010	<0.5	[NT]	[NT]	LCS	103%
Lead	mg/kg	1	SEM-010	<1	[NT]	[NT]	LCS	102%
Nickel	mg/kg	0.5	SEM-010	<0.5	[NT]	[NT]	LCS	102%
Zinc	mg/kg	0.5	SEM-010	<0.5	[NT]	[NT]	LCS	97%

QUALITY CONTROL Mercury Cold Vapor/Hg Analyser	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Matrix Spike % Recovery Duplicate + %RPD
Date Extracted (Mercury)				8/07/20 10	[NT]	[NT]	LCS	8/07/2010
Date Analysed (Mercury)				8/07/20 10	[NT]	[NT]	LCS	8/07/2010
Mercury	mg/kg	0.05	SEM-005	<0.05	[NT]	[NT]	LCS	108%

QUALITY CONTROL	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate	Spike Sm#	Matrix Spike % Recovery
VOCs by ZHE TCLP - 72 List						Base + Duplicate + %RPD		Duplicate + %RPD
pH of soil for fluid# determ.	pH units	0	AN101	[NT]	[NT]	[NT]	[NR]	[NR]
pH of soil for fluid # determ. (acid)	pH units	0	AN101	[NT]	[NT]	[NT]	[NR]	[NR]
Extraction fluid used	-		AN006	[NT]	[NT]	[NT]	[NR]	[NR]
pH of final Leachate	pH units	0	AN101	[NT]	[NT]	[NT]	[NR]	[NR]
Date extracted (VOCs)				08/07/2 010	[NT]	[NT]	LCS	08/07/2010
Date analysed (VOCs)				08/07/2 010	[NT]	[NT]	LCS	08/07/2010
Dichlorodifluoromethane (CFC-12)	µg/L	5	SEO-019/S EP-004	<5	[NT]	[NT]	[NR]	[NR]
Chloromethane	µg/L	5	SEO-019/S EP-004	<5	[NT]	[NT]	[NR]	[NR]
Vinyl Chloride (chloroethene)	µg/L	0.3	SEO-019/S EP-004	<0.3	[NT]	[NT]	[NR]	[NR]
Bromomethane	µg/L	10	SEO-019/S EP-004	<10	[NT]	[NT]	[NR]	[NR]



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QUALITY CONTROL	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate	Spike Sm#	Matrix Spike % Recovery
VOCs by ZHE TCLP - 72 List						Base + Duplicate + %RPD		Duplicate + %RPD
Chloroethane	µg/L	5	SEO-019/S EP-004	<5	[NT]	[NT]	[NR]	[NR]
Trichlorofluoromethane	µg/L	1	SEO-019/S EP-004	<1	[NT]	[NT]	[NR]	[NR]
Acetone (2-propanone)	µg/L	10	SEO-019/S EP-004	<10	[NT]	[NT]	[NR]	[NR]
1,1-Dichloroethene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	LCS	104%
Methyl lodine (iodomethane)	µg/L	5	SEO-019/S EP-004	<5	[NT]	[NT]	[NR]	[NR]
Acrylonitrile	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Methylene Chloride (DCM)	µg/L	5	SEO-019/S EP-004	<5	[NT]	[NT]	[NR]	[NR]
Allyl Chloride	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Carbon Disulphide	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
trans-1,2-Dichloroeth ene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Methyl-tert-butyl ether (MtBE)	µg/L	2	SEO-019/S EP-004	<2	[NT]	[NT]	[NR]	[NR]
1,1-Dichloroethane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
2-Butanone (MEK)	µg/L	10	SEO-019/S EP-004	<10	[NT]	[NT]	[NR]	[NR]
<i>cis</i> -1,2-Dichloroethen e	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Bromochloromethane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Chloroform	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	LCS	94%
2,2-Dichloropropane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,2-Dichloroethane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	LCS	105%
1,1,1-Trichloroethane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,1-Dichloropropene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Carbon tetrachloride	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Benzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	LCS	97%
Dibromomethane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]



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QUALITY CONTROL	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate	Spike Sm#	Matrix Spike % Recovery
VOCs by ZHE TCLP - 72 List						Base + Duplicate + %RPD		Duplicate + %RPD
1,2-Dichloropropane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Trichloroethene (TCE)	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	LCS	108%
2-Nitropropane	µg/L	100	SEO-019/S EP-004	<100	[NT]	[NT]	[NR]	[NR]
Bromodichloromethane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
<i>cis</i> -1,3-Dichloroprope ne	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
4-Methyl-2-Pentanone (MIBK)	µg/L	5	SEO-019/S EP-004	<5	[NT]	[NT]	[NR]	[NR]
<i>tran</i> s-1,3-Dichloropro pene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,1,2-Trichloroethane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Toluene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	LCS	112%
1,3-Dichloropropane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
2-Hexanone (MBK)	µg/L	5	SEO-019/S EP-004	<5	[NT]	[NT]	[NR]	[NR]
Dibromochloromethane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,2-Dibromoethane (EDB)	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Tetrachloroethene (PCE- perchloroethylen	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,1,1,2-Tetrachloroethan e	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Chlorobenzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	LCS	126%
Ethyl benzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	LCS	125%
Bromoform	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
<i>m/p</i> -Xylenes	µg/L	1	SEO-019/S EP-004	<1	[NT]	[NT]	LCS	132%
Cis-1,4-dichloro-2-buten e	µg/L	1	SEO-019/S EP-004	<1	[NT]	[NT]	[NR]	[NR]
Styrene (vinyl benzene)	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,1,2,2-Tetrachloroethan e	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
o-Xylene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	LCS	119%



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QUALITY CONTROL	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate	Spike Sm#	Matrix Spike % Recovery
VOCs by ZHE TCLP - 72 List						Base + Duplicate + %RPD		Duplicate + %RPD
1,2,3-Trichloropropane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Trans-1,4-dichloro-2-but ene	µg/L	1	SEO-019/S EP-004	<1	[NT]	[NT]	[NR]	[NR]
Isopropylbenzene (Cumene)	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Bromobenzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
n-Propylbenzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
2-Chlorotoluene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
4-Chlorotoluene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,3,5-Trimethylbenzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
tert-Butylbenzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,2,4-Trimethylbenzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
sec-Butylbenzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,3-Dichlorobenzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,4-Dichlorobenzene	µg/L	0.3	SEO-019/S EP-004	<0.3	[NT]	[NT]	[NR]	[NR]
p-lsopropyl toluene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,2-Dichlorobenzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
n-Butylbenzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,2-Dibromo-3-chloropro pane	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
1,2,4-Trichlorobenzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Naphthalene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Hexachlorobutadiene	µg/L	1	SEO-019/S EP-004	<1	[NT]	[NT]	[NR]	[NR]
1,2,3-Trichlorobenzene	µg/L	0.5	SEO-019/S EP-004	<0.5	[NT]	[NT]	[NR]	[NR]
Vinyl acetate	µg/L	10	SEO-019/S EP-004	<10	[NT]	[NT]	[NR]	[NR]
Dibromofluoromethane	%	0	SEO-019/S EP-004	90	[NT]	[NT]	LCS	88%



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REPORT NO: SE79619

QUALITY CONTROL	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate	Spike Sm#	Matrix Spike % Recovery
VOCs by ZHE TCLP - 72 List						Base + Duplicate + %RPD		Duplicate + %RPD
1,2-Dichloroethane-d4	%	0	SEO-019/S EP-004	83	[NT]	[NT]	LCS	86%
Toluene- <i>d8 Surrogate</i> 2	%	0	SEO-019/S EP-004	101	[NT]	[NT]	LCS	101%
4-Bromofluorobenzene Surrogate 3	%	0	SEO-019/S EP-004	85	[NT]	[NT]	LCS	85%

QUALITY CONTROL	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate	Spike Sm#	Matrix Spike % Recovery
PAHs in TCLP (USEPA 1311)						Base + Duplicate + %RPD		Duplicate + %RPD
pH of soil for fluid# determ.	pH units	0	AN101	[NT]	[NT]	[NT]	[NR]	[NR]
Extraction fluid used	-		AN006	[NT]	[NT]	[NT]	[NR]	[NR]
pH of soil for fluid # determ. (acid)	pH units	0	AN101	[NT]	[NT]	[NT]	[NR]	[NR]
pH of final Leachate	pH units	0	AN101	[NT]	[NT]	[NT]	[NR]	[NR]
Date Extracted				12/07/2 010	[NT]	[NT]	LCS	12/07/2010
Date Analysed				12/07/2 010	[NT]	[NT]	LCS	12/07/2010
Naphthalene	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	LCS	83%
2-Methylnaphthalene	µg/L	0.5	SEO-030	<0.5	[NT]	[NT]	[NR]	[NR]
1-Methylnaphthalene	µg/L	0.5	SEO-030	<0.5	[NT]	[NT]	[NR]	[NR]
Acenaphthylene	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	LCS	86%
Acenaphthene	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	LCS	95%
Fluorene	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	[NR]	[NR]
Phenanthrene	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	LCS	88%
Anthracene	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	LCS	95%
Fluoranthene	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	LCS	94%
Pyrene	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	LCS	96%
Benzo[a]anthracene	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	[NR]	[NR]
Chrysene	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	[NR]	[NR]
Benzo <i>[b,k</i>]fluoranthe ne	µg/L	1	SEO-030	<1.0	[NT]	[NT]	[NR]	[NR]
Benzo[a]pyrene	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	LCS	86%
Indeno[<i>123-cd</i>]pyren e	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	[NR]	[NR]
Dibenzo[<i>ah</i>]anthrace ne	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	[NR]	[NR]
Benzo[ghi]perylene	µg/L	0.5	SEO-030	<0.50	[NT]	[NT]	[NR]	[NR]
Total PAHs (sum)	µg/L	9	SEO-030	<9	[NT]	[NT]	[NR]	[NR]
Nitrobenzene-d5	%	0	SEO-030	96	[NT]	[NT]	LCS	87%



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REPORT NO: SE79619

QUALITY CONTROL	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate	Spike Sm#	Matrix Spike % Recovery
PAHs in TCLP (USEPA 1311)						Base + Duplicate + %RPD		Duplicate + %RPD
2-Fluorobiphenyl	%	0	SEO-030	84	[NT]	[NT]	LCS	83%
p -Terphenyl-d 14	%	0	SEO-030	95	[NT]	[NT]	LCS	92%

QUALITY CONTROL	UNITS	LOR	METHOD	Blank	Duplicate Sm#	Duplicate	Spike Sm#	Matrix Spike % Recovery
Metals in TCLP						Base + Duplicate + %RPD		Duplicate + %RPD
Date Extracted (Metals)				9/07/20 10	[NT]	[NT]	LCS	9/07/2010
Date Analysed (Metals)				9/07/20 10	[NT]	[NT]	LCS	9/07/2010
Date Extracted (Mercury)				9/07/20 10	[NT]	[NT]	LCS	9/07/2010
Date Analysed (Mercury)				9/07/20 10	[NT]	[NT]	LCS	9/07/2010
Arsenic	mg/L	0.05	SEM-010	<0.05	[NT]	[NT]	LCS	92%
Cadmium	mg/L	0.005	SEM-010	<0.005	[NT]	[NT]	LCS	97%
Chromium	mg/L	0.005	SEM-010	<0.005	[NT]	[NT]	LCS	100%
Copper	mg/L	0.01	SEM-010	<0.01	[NT]	[NT]	LCS	98%
Mercury	mg/L	0.0001	SEM-005	<0.000 1	[NT]	[NT]	LCS	102%
Nickel	mg/L	0.01	SEM-010	<0.010	[NT]	[NT]	LCS	96%
Lead	mg/L	0.02	SEM-010	<0.02	[NT]	[NT]	LCS	96%
Zinc	mg/L	0.01	SEM-010	<0.010	[NT]	[NT]	LCS	91%



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QUALITY CONTROL	UNITS	LOR	METHOD	Blank
Moisture				
Date Analysed (moisture)				[NT]
Moisture	%	1	AN002	<1



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Result Codes

 [INS]
 :
 Insufficient Sample for this test

 [NR]
 :
 Not Requested

 [NT]
 :
 Not tested

 [LOR]
 :
 Limit of reporting

 Report
 Comments

[RPD] : Relative Percentage Difference* : Not part of NATA Accreditation

[N/A] : Not Applicable

Samples analysed as received. Solid samples expressed on a dry weight basis.

Date Organics extraction commenced:

NATA Corporate Accreditation No. 2562, Site No 4354 Note: Test results are not corrected for recovery (excluding Air-toxics and Dioxins/Furans*) This document is issued by the Company subject to its General Conditions of Service (www.sgs.com/terms_and_conditions.htm). Attention is drawn to the limitations of liability, indemnification and jurisdictional issues established therein.

This document is to be treated as an original within the meaning of UCP 600. Any holder of this document is advised that information contained hereon reflects the Company's findings at the time of its intervention only and within the limits of client's instructions, if any. The Company's sole responsibility is to its Client and this document does not exonerate parties to a transaction from exercising all their rights and obligations under the transaction documents. Any unauthorized alteration, forgery or falsification of the content or appearance of this document is unlawful and offenders may be prosecuted to the fullest extent of the law.

Quality Control Protocol

Method Blank: An analyte free matrix to which all reagents are added in the same volume or proportions as used in sample processing. The method blank should be carried through the complete sample preparation and analytical procedure. A method blank is prepared every 20 samples.

Duplicate: A separate portion of a sample being analysed that is treated the same as the other samples in the batch. One duplicate is processed at least every 10 samples.

Surrogate Spike: An organic compound which is similar to the target analyte(s) in chemical composition and behavior in the analytical process, but which is not normally found in environmental samples. Surrogates are added to samples before extraction to monitor extraction efficiency and percent recovery in each sample.

Internal Standard: Added to all samples requiring analysis for organics (where relevant) or metals by ICP after the extraction/digestion process; the compounds/elements serve to give a standard of retention time and/or response, which is invariant from run-to-run with the instruments.

Laboratory Control Sample: A known matrix spiked with compound(s) representative of the target analytes. It is used to document laboratory performance. When the results of the matrix spike analysis indicates a potential problem due to the sample matrix itself, the LCS results are used to verify that the laboratory can perform the analysis in a clean matrix.

Matrix Spike: An aliquot of sample spiked with a known concentration of target analyte(s). The spiking occurs prior to sample preparation and analysis. A matrix spike is used to document the bias of a method in a given sample matrix.

Quality Acceptance Criteria

The QC criteria are subject to internal review according to the SGS QAQC plan and may be provided on request or alternatively can be found here: http://www.au.sgs.com/sgs-mp-au-env-qu-022-qa-qc-plan-en-09.pdf



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SAMPLE RECEIPT ADVICE (SRA)

9 July 2010

Client Details Requested By	: Sumi Dorairaj	Laboratory Details	5
Client	: JBS Environmental Pty Ltd	Laboratory :	SGS Environmental Services
Contact	: Sumi Dorairaj	Manager :	Edward Ibrahim
Address	: PO Box 940	Address :	Unit 16, 33 Maddox Street
	MASCOT NSW 1460		Alexandria NSW 2015
Email	: sdorairaj@jbsgroup.com.au	Email :	au.samplereceipt.sydney@sgs.com
Telephone	: 02 8338 1013	Telephone :	61 2 8594 0400
Facsimile	: 02 8338 1700	Facsimile :	61 2 8594 0499
Project	: 40913 - Macdonaldtown	Report No :	SE79619
Order Number	:	No. of Samples	1
Samples	: 1 Soil	Due Date :	12/07/2010
Date Instructions Received	: 5/07/2010		
Sample Receipt Date	: 5/7/10		
Samples received in good orde	er : YES	Samples received in correct container:	YES
Samples received without hea	adspace YES	Sufficient quantity supplied :	YES
Upon receipt sample temperatu	ure : Cool	Cooling Method :	Ice Pack
Sample containers provided by	y : Other Lab	Samples clearly Labelled :	YES
Turnaround time requested	: Standard	Completed documentation received :	YES

Samples will be held for 1 month for water samples and 3 months for soil samples from date of receipt of samples, unless otherwise instructed.

Comments

To the extent not inconsistent with the other provisions of this document and unless specifically agreed otherwise in writing by SGS, all SGS services are rendered in accordance with the applicable SGS General Conditions of Service accessible at http://www.sgs.com/terms_and_conditions.htm as at the date of this document. Attention is drawn to the limitations of liablility and to the clauses of indemnification.

The signed chain of custody will be returned to you with the original report.



SAMPLE RECEIPT ADVICE (SRA) - continued

Client	:	JBS Environmental Pty Ltd	Report No	:	SE79619
Project	:	40913 - Macdonaldtown			

Summary of Samples and Requested Analysis

The table below represents SGS Environmental Service's understanding and interpretation of the customer supplied sample request.

Please indicate ASAP if your request differs from these details.

Testing shall commence immediately as per this table, unless the customer intervenes with a correction prior to testing. Note that a small X in the table below indicates some testing has not been requested in the package.

Sample No.	Description	Metals Prep & Inorganics - All	VOCs in Soil-SGS Nat. 72 List	PAHs in Soil	Speciated Phenols in Soil	Metals in Soil by ICP-OES	Mercury Cold Vapor/Hg Analyser	VOCsZHE TCLP-SGS Nat. 72 List	PAHs in TCLP USEPA 1311	Metals in TCLP	Moisture
1	QC2	Х	Х	Х	Х	X	Х	Х	Х	Х	Х

Sample No.	Description
1	QC2

Signature: Envirolab Sample ID Email: sderaraje josgroup comay / tolavis & josgroup con av Client: Date & Time: Print Name: Phone: 8388 1011 Relinquished by (company): Address: Sampler: Project Mgr: Denaira 4126 22 57 20 J 5 2 G Form: 302 - Chain of Custody-Client, Issued 14/02/08, Version 3, Page 1 of 1. 082 128 O'Riorden St Mascol NSW 2020 JBS THO **UBS TP5** SUMI ORP QA1 lim Davis Client Sample ID Environmenta TPUS OF/10 N Draina 12.0m 153 Sample information 40-42 E n G 30 Fax: 02/07/10 Date sampled 4 8338 1700 **CHAIN OF CUSTODY - Client** Type of sample < in ENVIROLAB SERVICES Or chooser standard 1 day / 2 day / 3 day Signature: Client Project Name and Number: Date & Time: Print Name: Received by (company): Heavy Metals Envirolab Services Quote No. : surcharge applies Note: Inform lab in advance if urgent turnaround is required -PO No.: Date results required: < PAtls 40913 5 VOLS J speciated phenols 2-2 J Macdonalltown TCLP metals TCLP PAHIS seldu BELIG BIVEC Intaci Coller 5 TCLP VOCS TCLP phinols Ce-Cq BTEX õ **Tests Required** 517/108322 un HOLD Transported by: Hand delivered / courier Samples Received: Cool or Ambient (circle one) E-mail: tnotaras@envirolabservices.com.au Fax: **Envirolab** Services Temperature Recieved at: **Contact: Tania Notaras** Phone: 02 9958 5801 12 Ashley St, Chatswood, NSW, 2067 Ashestos 02 9958 5803 SION Sam YP1 108/6 180 0 80 polet Pack DBA les int Page No: please send to stes information about the (if applicable) sample as you can Provide as much Comments 2 cf 9

AU.SampleReceipt.Sydney (Sydney)

From:	Blackman, Daniel (Sydney)
Sent:	Friday, 9 July 2010 2:56 PM
To:	sdorairaj@jbsgroup.com.au
Cc:	AU.SampleReceipt.Sydney (Sydney)
Subject:	SE79619-1 - 40913 Macdonaldtown - Sample QC2
Attachments	: COC_SE79619.zip

Dear Sumi,

As instructed SGS will go ahead will TCLP PAHs. Note that we will not be able to test for TCLP speciated phenols due to the limited volume and 'rocky' nature of the soil sample.

Kind

Daniel Blackman Environmental Services Client Services Officer

SGS Australia Pty Ltd Unit 16, 33 Maddox St Alexandria, NSW, 2015 Phone: +61 (0)2 8594 0400 Fax: +61 (0)2 8594 0499 E-mail: daniel.blackman@sgs.com

SGS DataNet: View Results Online



Appendix G Tabulated Quality Control Results

Field Duplicates (soil) Filter: SDG in('42976')			SDG Field_ID Sampled_Date-Time	42976 JBS TP1 2/07/2010	42976 QA2 2/07/2010	RPD	42976 JBS TP1 2/07/2010	Interlab_D QC2 2/07/2010	RPD
Chem_Group BTEX	ChemName Benzene	Units mg/kg	EQL 0.5 (Primary): 0.1 (Interla	<0.5	<0.5	0	<0.5	<0.1	0
	Ethylbenzene Toluene Xylene (m & p)	µg/L mg/kg µg/L	1 (Primary): 0.5 (Interlab 0.5 (Primary): 0.1 (Interla 2 (Primary): 1 (Interlab)	<1.0 <0.5 <2.0	1.7 <0.5 2.7	52 0 30	<1.0 <0.5 <2.0	4.1 <0.1 9.0	122 0 127
	Xylene (o)	mg/kg	1 (Primary): 0.1 (Interlab	<1.0	<1.0	0	<1.0	0.1	0
Chlorinated Hydrocarbons	1,1,1,2-tetrachloroethane 1,1,1-trichloroethane 1,1,2,2-tetrachloroethane	mg/kg mg/kg mg/kg	1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0 <1.0	<1.0 <1.0 <1.0	0 0 0	<1.0 <1.0 <1.0	<0.1 <0.1 <0.1	0 0
	1,1,2-trichloroethane 1,1-dichloroethane	mg/kg mg/kg	1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0	<1.0 <1.0	0	<1.0 <1.0	<0.1 <0.1	0
	1,1-dichloroethene 1,1-dichloropropene 1,2,3-trichloropropane	mg/kg mg/kg mg/kg	1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0 <1.0	<1.0 <1.0 <1.0	0 0 0	<1.0 <1.0 <1.0	<0.1 <0.1 <0.1	0
	1,2-dibromo-3-chloropropane 1,2-dichloroethane	mg/kg μg/L	1 (Primary): 0.1 (Interlab 1 (Primary): 0.5 (Interlab	<1.0 <1.0	<1.0 <1.0	0	<1.0 <1.0	<0.1 <0.5	0
	1,2-dichloropropane 1,3-dichloropropane 2,2-dichloropropane	mg/kg mg/kg mg/kg	1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0 <1.0	<1.0 <1.0 <1.0	0 0 0	<1.0 <1.0 <1.0	<0.1 <0.1 <0.1	0 0 0
	Bromochloromethane Bromodichloromethane	mg/kg mg/kg	1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0	<1.0 <1.0	0	<1.0 <1.0	<0.1 <0.1	0
	Bromoform Carbon tetrachloride Chlorodibromomethane	mg/kg mg/kg mg/kg	1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0 <1.0	<1.0 <1.0 <1.0	0 0 0	<1.0 <1.0 <1.0	<0.1 <0.1 <0.1	0 0 0
	Chloroethane Chloroform Chloromethane	mg/kg mg/kg mg/kg	1 1 (Primary): 0.1 (Interlab	<1.0 <1.0 <1.0	<1.0 <1.0 <1.0	0 0 0	<1.0 <1.0 <1.0	<1.0 <0.1 <1.0	0 0 0
	cis-1,2-dichloroethene cis-1,3-dichloropropene	mg/kg µg/L	1 (Primary): 0.1 (Interlab 1 (Primary): 0.5 (Interlab	<1.0 <1.0 <1.0	<1.0 <1.0 <1.0	0	<1.0 <1.0 <1.0	<0.1 <0.5	0
	Dibromomethane Hexachlorobutadiene Trichloroethene	mg/kg mg/kg mg/kg	1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0 <1.0	<1.0 <1.0 <1.0	0 0 0	<1.0 <1.0 <1.0	<0.1 <0.1 <0.1	0
	Tetrachloroethene trans-1,2-dichloroethene	mg/kg mg/kg	1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0	<1.0 <1.0	0	<1.0 <1.0	<0.1 <0.1	0
	trans-1,3-dichloropropene Vinyl chloride	µg/L mg/kg	1 (Primary): 0.5 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0	<1.0 <1.0	0	<1.0 <1.0	<0.5 <0.1	0
Halogenated Benzenes	1,2,3-trichlorobenzene 1,2,4-trichlorobenzene	mg/kg mg/kg	1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0	<1.0 <1.0	0	<1.0 <1.0	<0.1 <0.1	0
	1,2-dichlorobenzene 1,3-dichlorobenzene 1,4-dichlorobenzene	mg/kg mg/kg µg/L	1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab 1 (Primary): 0.3 (Interlab	<1.0 <1.0 <1.0	<1.0 <1.0 <1.0	0 0 0	<1.0 <1.0 <1.0	<0.1 <0.1 <0.3	0 0 0
	2-chlorotoluene 4-chlorotoluene	mg/kg µg/L	1 (Primary): 0.1 (Interlab 1 (Primary): 0.5 (Interlab	<1.0 <1.0	<1.0 <1.0	0	<1.0 <1.0	<0.1 <0.5	0
	Bromobenzene Chlorobenzene	mg/kg mg/kg	1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0	<1.0 <1.0	0	<1.0 <1.0	<0.1 <0.1	0
Halogenated Hydrocarbons	1,2-dibromoethane Bromomethane	µg/L mg/kg	1 (Primary): 0.5 (Interlab	<1.0 <1.0	<1.0 <1.0	0	<1.0 <1.0	<0.5 <1.0	0
	Dichlorodifluoromethane Trichlorofluoromethane	mg/kg mg/kg	1 1	<1.0 <1.0	<1.0 <1.0	0	<1.0 <1.0	<1.0 <1.0	0
Halogenated Phenols	2,3,4,6-tetrachlorophenol 2,4,5-trichlorophenol	mg/kg		<1.0 <1.0	<1.0 <1.0	0	<1.0 <1.0	<0.5 <0.5	0
	2,4,6-trichlorophenol 2,4-dichlorophenol 2,6-dichlorophenol	mg/kg mg/kg mg/kg		<1.0 <1.0 <1.0	<1.0 <1.0 <1.0	0 0 0	<1.0 <1.0 <1.0	<0.5 <0.5 <0.5	0 0 0
	2-chlorophenol Pentachlorophenol		1 (Primary): 0.5 (Interlab 10 (Primary): 0.5 (Interla	<1.0 <10.0	<1.0 <10.0	0	<1.0 <10.0	<0.5 <0.5	0
Inorganics	Moisture	%	0.1 (Primary): 1 (Interlab	13.0	17.0	27	13.0	17.0	27
Lead (leached)	Lead Lead (Filtered)	mg/kg mg/l	0.03	220.0 <0.03	280.0	24 80	<0.03	290.0	27
MAH	1,2,4-trimethylbenzene	mg/kg	1 (Primary): 0.1 (Interlab	<1.0	<1.0	0	<1.0	0.3	0
	1,3,5-trimethylbenzene Isopropylbenzene n-butylbenzene	mg/kg mg/kg µg/L	1 (Primary): 0.1 (Interlab 1 (Primary): 0.1 (Interlab 1 (Primary): 0.5 (Interlab	<1.0 <1.0 <1.0	<1.0 <1.0 <1.0	0 0 0	<1.0 <1.0 <1.0	0.1 <0.1 <0.5	0 0 0
	n-propylbenzene p-isopropyltoluene	μg/L mg/kg	1 (Primary): 0.5 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0	<1.0 <1.0	0	<1.0 <1.0	<0.5 <0.1	0
	sec-butylbenzene Styrene tert-butylbenzene	mg/kg µg/L mg/kg	1 (Primary): 0.1 (Interlab 1 (Primary): 0.5 (Interlab 1 (Primary): 0.1 (Interlab	<1.0 <1.0 <1.0	<1.0 <1.0 <1.0	0 0 0	<1.0 <1.0 <1.0	<0.1 <0.5 <0.1	0 0 0
Metals	Arsenic	mg/kg	4 (Primary): 3 (Interlab)	30.0	30.0	0	30.0	45.0	40
	Cadmium Chromium (III+VI) Copper	mg/kg	0.5 (Primary): 0.3 (Interla 1 (Primary): 0.3 (Interlab 1 (Primary): 0.5 (Interlab	1.1 26.0 230.0	1.1 21.0 260.0	0 21 12	1.1 26.0 230.0	2.1 23.0 380.0	63 12 49
	Mercury Nickel	mg/kg mg/kg	0.1 (Primary): 0.05 (Inter 1 (Primary): 0.5 (Interlab	0.3 20.0	0.4 24.0	29 18	0.3 20.0	0.26 24.0	14 18
Metals (leached)	Zinc Arsenic (Filtered)	mg/kg mg/l	1 (Primary): 0.5 (Interlab	260.0 <0.05	330.0 <0.05	24 0	260.0 <0.05	370.0	35
	Cadmium (Filtered) Chromium (III+VI) (Filtered)	mg/l mg/l	0.01 0.01	<0.01 <0.01	<0.01 <0.01	0	<0.01 <0.01		
	Copper (Filtered) Mercury (Filtered) Nickel (Filtered)	mg/l mg/l mg/l	0.01 0.0005 0.02	0.1 <0.0005 0.02	0.1 <0.0005 0.05	0 0 86	0.1 <0.0005 0.02		
DAH/Dhopolo	Zinc (Filtered)	mg/l	0.02	1.6	1.8 <1.0	12	1.6	<0.5	0
PAH/Phenols	2,4-dimethylphenol 2,4-dinitrophenol 2-methylphenol	mg/kg mg/kg mg/kg	1 (Primary): 0.5 (Interlab 10 (Primary): 0.5 (Interla 1 (Primary): 0.5 (Interlab	<1.0 <10.0 <1.0	<10.0 <10.0 <1.0	0 0 0	<1.0 <10.0 <1.0	<0.5 <0.5 <0.5	0 0 0
	2-nitrophenol 4,6-Dinitro-2-methylphenol 4-methylphenol	mg/kg mg/kg mg/kg	1 (Primary): 0.5 (Interlab 10	<1.0 <10.0 <2.0	<1.0 <10.0 <2.0	0 0 0	<1.0 <10.0 <2.0	<0.5	0
	4-nitrophenol Acenaphthene	mg/kg mg/kg	0.1	<10.0 0.1	<10.0 0.2	0 67	<10.0 0.1	<0.5 0.23	0 79
	Acenaphthylene Anthracene Benz(a)anthracene	mg/kg mg/kg mg/kg	0.1	0.7 0.8 2.6	1.7 2.1 10.0	83 90 117	0.7 0.8 2.6	2.4 2.6 11.0	110 100
	Benzo(a) pyrene Benzo(b)&(k)fluoranthene	mg/kg mg/kg	0.05 0.2	3.0 3.8	11.0 14.0	114 115	3.0 3.8	6.7 12.0	76 104
	Benzo(g,h,i)perylene Chrysene Dibenz(a,h)anthracene	mg/kg mg/kg mg/kg	0.1	1.1 2.6 0.3	4.4 9.1 1.2	120 111 120	1.1 2.6 0.3	4.6 6.3 1.2	12: 83 12(
	Fluoranthene Fluorene	mg/kg mg/kg	0.1 0.1	4.4 0.5	15.0 0.7	109 33	4.4 0.5	12.0 0.92	93 59
	Indeno(1,2,3-c,d)pyrene Naphthalene Phenanthrene	mg/kg mg/kg mg/kg	0.1 0.1	1.3 0.6 3.4	4.6 2.8 7.9	112 129 80	1.3 0.6 3.4	4.3 3.1 - 5.2 5.7	107 159 51
	Phenol Pyrene		1 (Primary): 0.5 (Interlab	<1.0 4.9	<1.0 20.0	0 121	<1.0 4.9	<0.5 13.0	0 91
PAH/Phenols (leached)	Acenaphthene (Filtered) Acenaphthylene (Filtered)	μg/l μg/l	1	<1.0 <1.0	1.0 <1.0	0	<1.0 <1.0		E
	Anthracene (Filtered) Benz(a)anthracene (Filtered) Benzo(a) pyrene (Filtered)	µg/l µg/l µg/l	1	<1.0 <1.0 <1.0	1.0 <1.0 <1.0	0 0 0	<1.0 <1.0 <1.0		
	Benzo(b)&(k)fluoranthene (Filtered) Benzo(g,h,i)perylene (Filtered)	μg/l μg/l	1 2 1	<2.0 <1.0	<2.0 <1.0	0	<2.0 <1.0		F
	Chrysene (Filtered) Dibenz(a,h)anthracene (Filtered) Fluoranthene (Filtered)	μg/l μg/l μg/l	1 1 1	<1.0 <1.0 <1.0	<1.0 <1.0 <1.0	0 0 0	<1.0 <1.0 <1.0		
	Fluorene (Filtered) Indeno(1,2,3-c,d)pyrene (Filtered)	μg/l μg/l	1 1	<1.0 <1.0	1.0 <1.0	0	<1.0 <1.0		E
	Naphthalene (Filtered) Phenanthrene (Filtered) Phenolics Total	µg/l µg/l µg/l	1 1 50	<1.0 <1.0 <50.0	30.0 1.0 <50.0	187 0 0	<1.0 <1.0 <50.0		
	Pyrene (Filtered)	µg/i µg/l	1	<1.0	<50.0	0	<1.0		

Field Duplicates (soil)	SDG	42976	42976		42976	Interlab_D	
Filter: SDG in('42976')	Field_ID	JBS TP1	QA2	RPD	JBS TP1	QC2	RPD
	Sampled_Date-Time	2/07/2010	2/07/2010		2/07/2010	2/07/2010	

**High RPDs are in bold (Acceptable RPDs for each EQL multiplier range are: 50 (4-10 x EQL); 50 (10-30 x EQL); 50 (> 30 x EQL))
***Interlab Duplicates are matched on a per compound basis as methods vary between laboratories. Any methods in the row header relate to those used in the primary laboratories.

Field Blanks (water) Filter: SDG in('42976')			SDG Field_ID Sampled_Date-Time Sample_Type	42976 Rinsate 2/07/2010 Rinsate
Chem_Group	ChemName	Units	EQL	
BTEX	Benzene	µg/L	1	<1
BIEX	Ethylbenzene	µg/L	1	<1
	Toluene	µg/L	1	<1
	Xylene (m & p)	µg/L	2	<2
	Xylene (o)	µg/∟ µg/L	1	<1
		µy/∟	1	<u> </u>
Chlorinated Hydrocarbons	1,1,1,2-tetrachloroethane	µg/L	1	<1
Chionnaled Hydrocarbons	1,1,1-trichloroethane	µg/∟ µg/L	1	<1
	1,1,2,2-tetrachloroethane	µg/∟ µg/L	1	<1
	1,1,2-trichloroethane			<1
		µg/L	1	<1
	1,1-dichloroethane 1,1-dichloroethene	µg/L	1	
	,	µg/L		<1
	1,1-dichloropropene	µg/L	1	<1
	1,2,3-trichloropropane	µg/L	1	<1
	1,2-dibromo-3-chloropropane	µg/L	1	<1
	1,2-dichloroethane	µg/L	1	<1
	1,2-dichloropropane	µg/L	1	<1
	1,3-dichloropropane	µg/L	1	<1
	2,2-dichloropropane	µg/L	1	<1
	Bromochloromethane	µg/L	1	<1
	Bromodichloromethane	µg/L	1	<1
	Bromoform	µg/L	1	<1
	Carbon tetrachloride	μg/L	1	<1
	Chlorodibromomethane	µg/L	1	<1
	Chloroethane	µg/L	10	<10
	Chloroform	µg/L	1	<1
	Chloromethane	µg/L	10	<10
	cis-1,2-dichloroethene	µg/L	1	<1
	cis-1,3-dichloropropene	μg/L	1	<1
	Dibromomethane	µg/L	1	<1
	Hexachlorobutadiene	µg/L	1	<1
	Trichloroethene	µg/L	1	<1
	Tetrachloroethene	µg/L	1	<1
	trans-1,2-dichloroethene	µg/L	1	<1
	trans-1,3-dichloropropene	µg/L	1	<1
	Vinyl chloride	µg/L	10	<10
		P9/L		
Halogenated Benzenes	1,2,3-trichlorobenzene	ua/I	1	<1
alogenaleu Denzenes		µg/L	1	-
	1,2,4-trichlorobenzene 1,2-dichlorobenzene	µg/L	1	<1 <1
		µg/L		
	1,3-dichlorobenzene	µg/L	1	<1
	1,4-dichlorobenzene	µg/L	1	<1
	2-chlorotoluene	µg/L	1	<1
	4-chlorotoluene	µg/L	1	<1
	Bromobenzene	µg/L	1	<1
	Chlorobenzene	µg/L	1	<1
Halogenated Hydrocarbons	1,2-dibromoethane	µg/L	1	<1
	Bromomethane	µg/L	10	<10
	Dichlorodifluoromethane	µg/L	10	<10
	Trichlorofluoromethane	µg/L	10	<10
Halogenated Phenols	2,3,4,6-tetrachlorophenol	µg/L	10	<10
	2,4,5-trichlorophenol	µg/L	10	<10
	2,4,6-trichlorophenol	µg/L	10	<10
	2,4-dichlorophenol	µg/L	10	<10
	2,6-dichlorophenol	µg/L	10	<10
	2-chlorophenol	µg/L	10	<10
	Dentechlerenhenel	<u>µg</u> ,∟	100	<100

Lead	Lead (Filtered)	mg/l	0.03	<0.03
MAH	1,2,4-trimethylbenzene	µg/L	1	<1
	1,3,5-trimethylbenzene	µg/L	1	<1
	Isopropylbenzene	µg/L	1	<1
	n-butylbenzene	µg/L	1	<1
	n-propylbenzene	μg/L	1	<1
	p-isopropyltoluene	µg/L	1	<1
	sec-butylbenzene	µg/L	1	<1
	Styrene	µg/L	1	<1
	tert-butylbenzene	µg/L	1	<1
Metals	Arsenic (Filtered)	mg/l	0.05	<0.05
	Cadmium (Filtered)	mg/l	0.01	<0.01
	Chromium (III+VI) (Filtered)	mg/l	0.01	<0.01
	Copper (Filtered)	mg/l	0.01	<0.01
	Mercury (Filtered)	mg/l	0.0005	< 0.0005

µg/L

100

<100

Pentachlorophenol

Field Blanks (water)			SDG	42976
Filter: SDG in('42976')			Field_ID	Rinsate
, , , , , , , , , , , , , , , , , , ,			Sampled_Date-Time	2/07/2010
			Sample_Type	Rinsate
	Nickel (Filtered)	mg/l	0.02	<0.02
	Zinc (Filtered)	mg/l	0.02	<0.02
PAH/Phenols	2,4-dimethylphenol	µg/L	10	<10
	2,4-dinitrophenol	mg/l	0.1	<0.1
	2-methylphenol	µg/L	10	<10
	2-nitrophenol	µg/L	10	<10
	4,6-Dinitro-2-methylphenol	µg/L	100	<100
	4-methylphenol	mg/l	0.02	<0.02
	4-nitrophenol	µg/L	100	<100
	Acenaphthene	μg/L	1	<1
	Acenaphthylene	µg/L	1	<1
	Anthracene	μg/L	1	<1
	Benz(a)anthracene	µg/L	1	<1
	Benzo(a) pyrene	µg/L	1	<1
	Benzo(b)&(k)fluoranthene	µg/L	2	<2
	Benzo(g,h,i)perylene	µg/L	1	<1
	Chrysene	µg/L	1	<1
	Dibenz(a,h)anthracene	µg/L	1	<1
	Fluoranthene	µg/L	1	<1
	Fluorene	µg/L	1	<1
	Indeno(1,2,3-c,d)pyrene	µg/L	1	<1
	Naphthalene	µg/L	1	<1
	Phenanthrene	µg/L	1	<1
	Phenol	µg/L	10	<10
	Pyrene	µg/L	1	<1
Solvents	Cycloboxano	ma/l	0.001	<0.001
SUIVEIIIS	Cyclohexane	mg/l	0.001	<u>\0.001</u>



Appendix H Test Pit Logs



Project: Macdonaldtown Gasworks

Project No.: 40913

Client: Incoll Management

Project Manager: Sumi Dorairaj

Total Hole Depth: 1.0 m Eastings: -Northings: -Date: 02/07/2010 Operator and Co.: Anthony Colluso Excavation Method: Backhoe Log By: Sumi Dorairaj Excavation Width: 450 mm

	SUE	BSURFAC	CE PROFILE	SAMPLE			
Depth	Visual		Description	Number	Condition	PID (ppm)	Observations
-0.0		coke and damp, bal (irregular, FILL Silty Clay, red ironsto compacted	est pit at 1.0 m. Target	JBS TP1-0.3-0.4+QA/C2	D		Bulk sample collected
- - - - - - - - - - - - - - - - - - -							
- 3.0 - 3.0 							
- - - - -	Method		Sample Condition				
Sample MethodSample ConditionHA - Hand AugerU - undisturbed tube sampleSFA - Solid Flight AugerD - disturbed sampleHFA - Hollow Flight AugerCS - core samplePT - Push TubingImage: CS - core sample		ple					



Project: Macdonaldtown Gasworks

Project No.: 40913

Client: Incoll Management

Project Manager: Sumi Dorairaj

Total Hole Depth: 1.8 m Eastings: -Northings: -Date: 02/07/2010 Operator and Co.: Anthony Colluso Excavation Method: Backhoe Log By: Sumi Dorairaj Excavation Width: 450 mm

SUBSURFACE PROFILE				SAMPLE				
Depth	Visual		Description	Number	Condition	PID (ppm)	Observations	
-0.0		black, col (black, and 'coal tar' lii boulders, at 0.15m, mm x 450 FILL Silty Clay, compacted	Ground Surface elly Sand, dark brown to ke and slag inclusions gular, hard, fine to course), ke odour, large 'coke' dry to damp, asphalt layer steel plate below (1500 mm x 6 mm thick) orange mottled grey, well d with red streaks, slight ke odour, damp	JBS TP2-0.4 - 0.5 JBS TP2-0.8 - 0.9 JBS TP2-1.2 - 1.3	D			
- - - - - - - - - - - - - - - - - - -		Seepage into pit (brown, water, localised to 0.9 - 0.95m) FILL Grades to brown mottled orange, wet, slight 'coal tar' like odour Fill Silty Clay, dark grey with black, oily sheen / ooze, strong 'coal' like odour, wet End of test pit at 1.8 m. Refusal possibly on concrete.		JBS TP2-1.4 - 1.5	D			
	Method		Sample Condition					
HA - Hand AugerU - undisturbed tube sampleSFA - Solid Flight AugerD - disturbed sampleHFA - Hollow Flight AugerCS - core samplePT - Push Tubing			ple					



Project: Macdonaldtown Gasworks

Project No.: 40913

Client: Incoll Management

Project Manager: Sumi Dorairaj

Total Hole Depth: 4.3 m Eastings: -Northings: -Date: 02/07/2010 Operator and Co.: Anthony Colluso Excavation Method: Backhoe Log By: Sumi Dorairaj Excavation Width: 450 mm

SUBSURFACE PROFILE				SAMPLE				
Depth	Visual	Description		Number	Condition	PID (ppm)	Observations	
		Ground Surface FILL Silty Sandy Gravel, dark black coarse, medium grain, damp, heterogeneous with rootlets and road base gravels FILL Silty Clay, red orange mottled with grey and yellow, medium plasticity, damp - wet, heterogeneous with slight coal tar odour					Grass cover	
- - - - - - - - - - - - - - - -		FILL Silty Clay, dark brown, low plasticity, wet, heterogeneous with coke gravels and tar, strong coal tar odours and sheen FILL Silty Clay, yellow red with grey		JBS TP3-1.3-1.7	D		Bulk sample collected	
- - - - - - - - - - - - - -		FILL						
-4.0 - - -	As Above, very strong coal tar odours, wet, seepage of black ooz throughout End of test pit at 4.3 m. Equipment refusal.		et, seepage of black ooze t est pit at 4.3 m.	JBS TP3-4.0-4.2	U		Bulk and Bulk control samples collected	
Sample	Sample Method Sample Condition							
HA - Hand Auger SFA - Solid Flight Auger HFA - Hollow Flight Auger PT - Push Tubing			ble					



Project: Macdonaldtown Gasworks

Project No.: 40913

Client: Incoll Management

Project Manager: Sumi Dorairaj

Total Hole Depth: 2.1 m Eastings: -Northings: -Date: 02/07/2010 Operator and Co.: Anthony Colluso Excavation Method: Backhoe Log By: Tim Davis Excavation Width: 450 mm

SUBSURFACE PROFILE				SAMPLE				
Depth	Visual		Description	Number	Condition	PID (ppm)	Observations	
-0.0	×××××	FILL	Ground Surface				Grass cover	
- - - - - - - - - - - - - - - - - - -		Silty Sand medium g heterogen sandstone ACM fragr glass fragr 0.5 m FILL Silty Sand	, dark brown, fine to rain, dry to damp, eous with inclusions of e pieces, few suspected ments, timber, ash/slag, ments, slight PAH odour at with crushed sandstone	JBSTP4-0.4-0.5 + QA1 JBSTP4-0.9-1.0	D		No sample collected	
		FILL Silty Clay, mottles, m heterogen odour FILL	grey white mix / red orange with grey hedium plasticity, damp, eous, very slight PAH / grey brown, medium	JBSTP4-1.6-1.7 JBSTP4-1.9-2.0	D			
-2.0 - - - - - -		Plasticity, damp, heterogeneous, inclusion of shale pieces to sheets, few plastic pieces End of borehole at 2.1 m. Target depth reached.						
- -3.0 - - - -								
- - - 4.0 - - - -								
Sample	Sample Method Sample Condition							
HA - Hand AugerU - undisturbed tube sanSFA - Solid Flight AugerD - disturbed sampleHFA - Hollow Flight AugerCS - core samplePT - Push TubingImage: CS - core sample			ple					



Project: Macdonaldtown Gasworks

Project No.: 40913

Client: Incoll Management

Project Manager: Sumi Dorairaj

Total Hole Depth: 2.1 m Eastings: -Northings: -Date: 02/07/2010 Operator and Co.: Anthony Colluso Excavation Method: Backhoe Log By: Tim Davis Excavation Width: 450 mm

SUBSURFACE PROFILE				SAMPLE				
Depth	Visual		Description	Number	Condition	PID (ppm)	Observations	
-0.0		Ground Surface FILL Grass cover. Roadbase and bitumen mix, some sand, heterogeneous with inclusion of ash/slag FILL Silty Clay, orange brown to yellow, medium plasticity, damp, heterogeneous, inclusions of roadbase gravels, moderate to strong coal tar odours FILL As above FILL As above, strong coal tar odours FILL As above, strong coal tar odours End of test pit at 2.10m. End of investigation.		JBS TP5-0.4-0.5 JBS TP5-1.0-1.1 JBS TP5-1.5-1.6 JBS TP5-1.9-2.0	D			
Sample	Sample Method Sample Condition							
Sample wearloaSample contationHA - Hand AugerU - undisturbed tube sampleSFA - Solid Flight AugerD - disturbed sampleHFA - Hollow Flight AugerCS - core samplePT - Push TubingImage: Contation				ble				



Appendix I Pump Test Field Sheets

Field Reporting Cover Sheet



	PROJECT NAME: Macdonaldtown Siding PROJECT NO: 400	113
	FIELD DATES: Gth - 9th /7/2010 JBS PM: S. DORA	irat
	FIELD STAFF: T. DAVIS	·····
	WEATHER: RAIN	
	PROPOSED WORKS:	
	SUMMARY OF FIELDWORK: - commission of trial water treatment	
	- water pumped from wells MWDHS & MW373	
	- total volume ~4000L as described in field notes mixed	with
	4000L on typ water to increase pump volume	······································
\bigcirc	IMPORTANT OUTCOMES TO NOTE: Dutalogging triels to be conducted	in coming
	wæK	
	COC SENT? Yes DATE SAMPLES SENT: 9/7/10	
	PRIMARY LAB: ENVIROLAS SECONDARY LAB: NO	
ļ		
	POST-FIELDWORK CHECKLIST	
Jan Star	COC checked with PM	
	Site map updated and attached	
	Field data sheets attached (bore logs, well monitoring sheets)	
	Calibration and decontamination sheets attached	
	Field notes attached	
	Photos downloaded to file	
	Electronic bore logs completed and checked	
	OUTSTANDING ACTIONS:	

ENVIE CLIENT/SITE: DATE: **PARTICIPANTS:** PAGE: OF: MW423 Depth water - 1.020 to 40 Depth after pumping - 2.290 ater Total Depty - 4.344 4 pump rate ="41/min 1 . O-Imin Ø 2-3 min = 2.7 L/min 4-5 mins = 1.8 L/min = 1.86 (min 6-7 mins 8-9 min = 1.7 L/min -× Turn pump off after 10ming @ 10.37 × Renove datalogges after the @ 11.37 IMSO Forms020 - JBS Pad

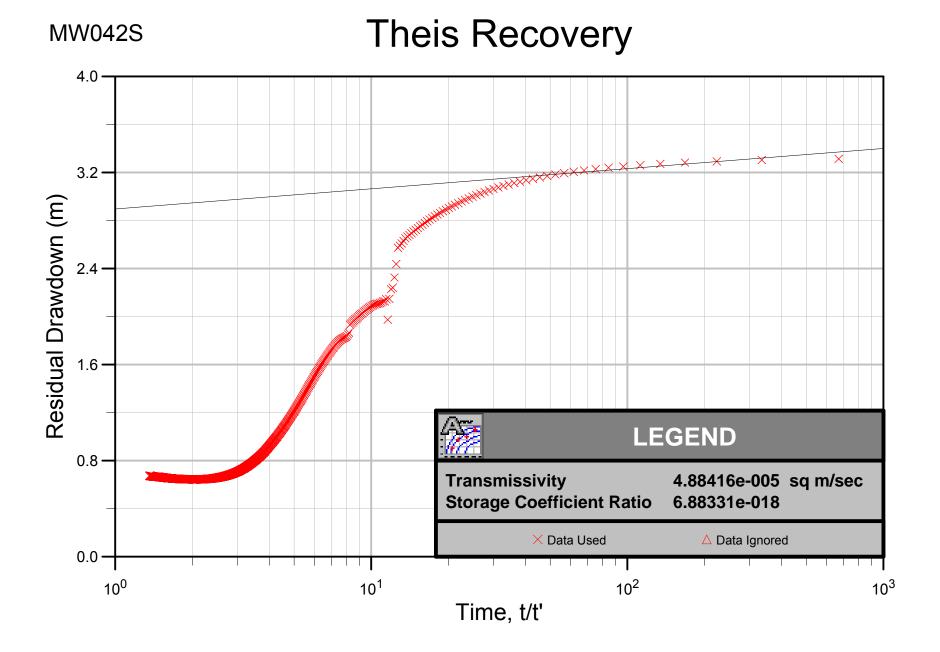
128 O'Riordan St Mascot NSW 2020 Tel +61 2 8338 1011 Fax +61 2 8338 1700 www.jbsgroup.com.au

CLIENT/SITE: DATE: 11/8/10 **PARTICIPANTS:** PAGE: OF: MW645 to writer before punping 1.359 Deepth 4.263 Depth Total - 1.421 after 6 13 mm writer after alter PUNCTIN Depth RMA Start proping 8.504 0-1mm = 3.9 L/min 2-3mm 4.2L/mm -----4-5min = 4.4 L/min 6-7 mins = 4.3 L/min 8-9nms = 4.2 L/min removed PVING ويناحك 12 mins ditalogger 221~0 9.5000 KO \$ γ () 1 IMSO Forms020 - JBS Pad

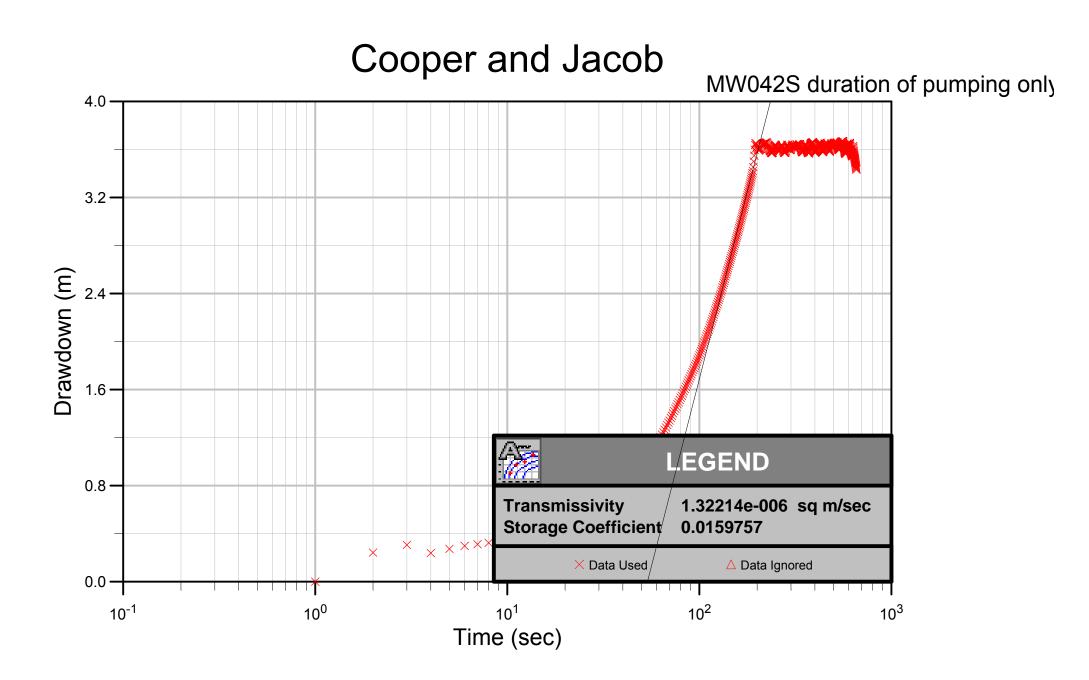
128 O'Riordan St Mascot NSW 2020 Tel +61 2 8338 1011 Fax +61 2 8338 1700 www.jbsgroup.com.au

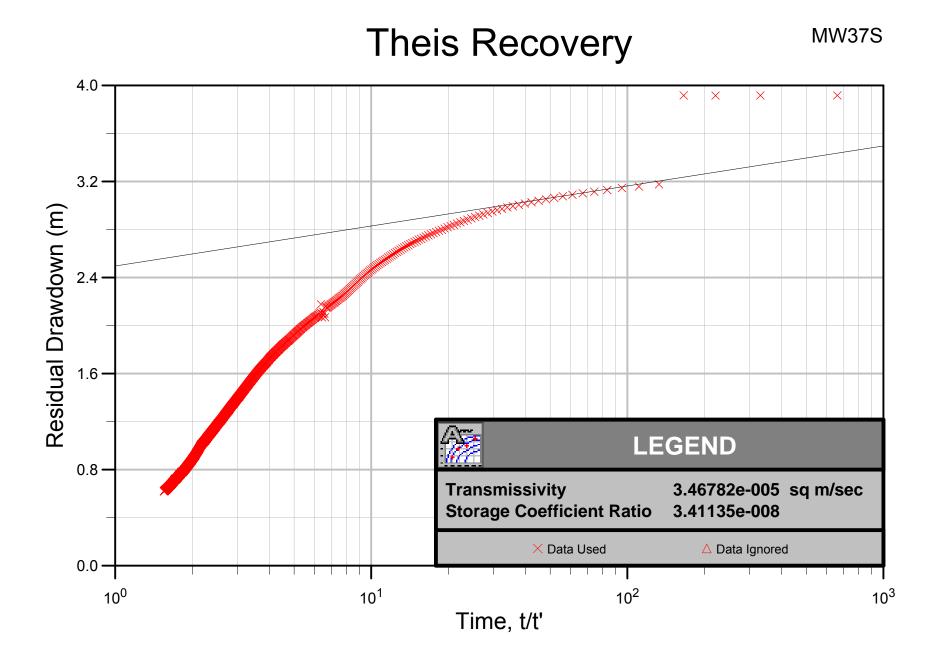
	-	
CLIENT/SITE:	DATE: \	18/10
PARTICIPANTS:	PAGE:	OF:
MWOGS		·
Depth to water	- 2.081	
Depth to water after pumpin	4-3.139	
rotal Depth	- 4.374 4	
Pump Rate		
$\frac{0-1\min}{2-3\min} = \frac{3L}{\min}$		
$\frac{2-5\min = 2L/\min}{4-5\min = 1.1L/\min}$		
G-7min = 0/min		
8-qmins = OL/min		
<u> </u>		
× pump removed after 10.		
× pump removed after 10 m × Datalogger removed rufter		

00 www.jbsgrou om.au

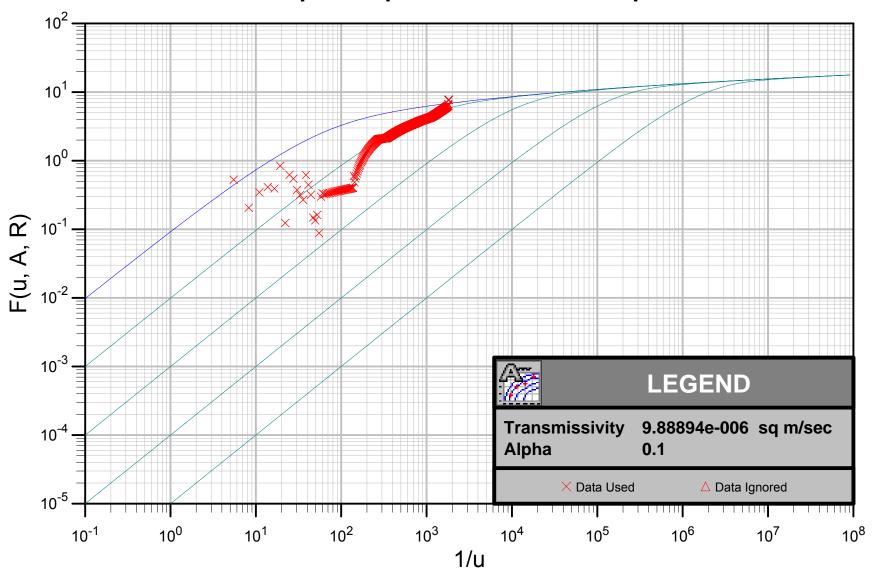


Papadopulos and Cooper MW042S duration of pumping only 10² 10¹ 10⁰ 10⁻² -10⁻³ LEGEND Transmissivity 9.69907e-006 sq m/sec 10⁻⁴ Alpha 0.1 × Data Used △ Data Ignored 10⁻⁵ 10³ 10⁻¹ 10⁰ 10¹ 10² 10⁴ 10⁵ 10⁶ 10⁷ 10⁸ 1/u



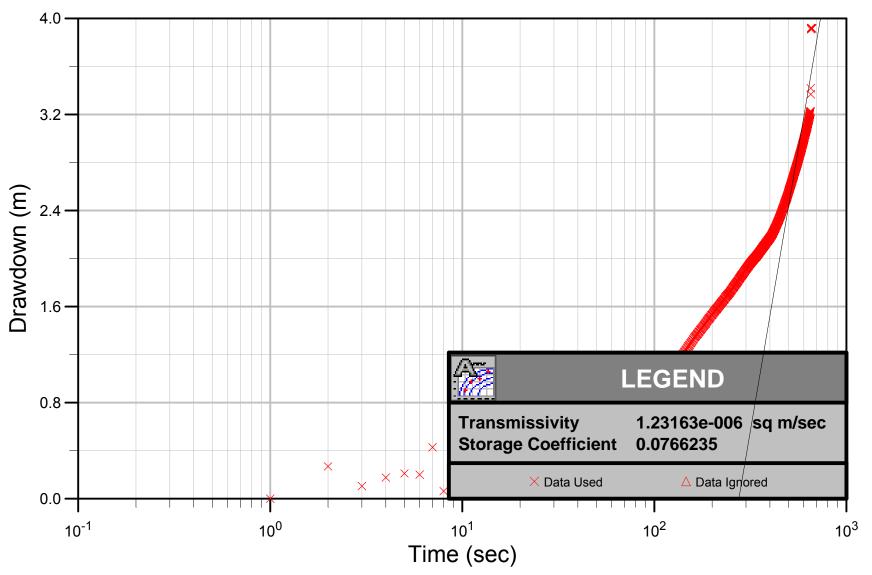


MW37S Duration of pumping onl





MW37S Duration of pumping onl





Appendix J

Results of Geotechnical Testing and Benchscale Stabilisation Trial



JBS Environmental Pty Ltd 128 O'Riordan Street Mascot, NSW 2028 Email: SDorairaj@jbsenvironmental.com.au

16 August, 2010

Attention: Sumi Dorairaj

RE: MACDONALDTOWN GASWORKS TREATMENT TRIAL

Dear Sumi,

Enviropacific Services Pty Ltd (EPS) is pleased to provide JBS Environmental Pty Ltd (JBS) with this report on the above works. Client satisfaction is the highest priority on all our projects and accordingly, we pride ourselves on our ability to offer innovative and flexible project delivery approaches to ensure client needs are satisfied. We also place the highest importance on OHS&R, environmental management, integrity and compliance with relevant laws and regulations. This commitment to quality is evidenced through obtaining third party accreditation of our Management System to ISO9001-2008 and accreditation for OHS&R and Environmental System to AS4801 and ISO14001 respectively.

Definitions

• ASLP – Leachable Concentration assessed by the Australian Standard Leaching Procedure (AS 4439.3-1997 Wastes, Sediments and Contaminated Soils - Preparation of Leachates - Bottle Leaching Procedures)

- BaP Benzo(a)pyrene
- DECCW NSW Department of Environment, Climate Change and Water
- EPS Enviropacific Services Pty Ltd
- General IA DECC General Approval for the Immobilisation of Contaminants in Waste, # 2005/14 Stabilisation of Coal Tar Contaminated Soil
- IA –Immobilisation Approval
- JBS JBS Environmental Pty Ltd
- PAH Polycyclic Aromatic Hydrocarbon
- SCC Specific Contaminant Concentration
- TCLP Leachable Concentration assessed by the Toxicity Characteristic Leaching Procedure

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- TPH Total Petroleum Hydrocarbon
- UCS Unconfined Compressive Strength measured by NSW RTA Test Method T131

• Waste Classification Guidelines – Refers to Table 2 of DECCW Waste Classification Guidelines, Part 1: Classifying Waste (2009)

Scope of work

Enviropacific was engaged by JBS to conduct an immobilisation treatment trial using Portland cement to immobilise three different materials originating from the former Macdonaldtown Gasworks.

The treatment trial included the following tasks:

1. Subsamples of each material were blended and homogenised in accordance with the flowchart shown below in Figure 1.

2. Each material was prepared in accordance with NSW RTA Test Method T131.

3. Three samples from each prepared material (total 9) were collected for pre-treatment chemical analysis. Each sample was analysed for the following:

- As, Cd, Cr, Pb, Hg, Ni (SCC and TCLP)
- Benzo(a)pyrene and PAHs (SCC and TCLP)
- Phenols (SCC and TCLP)
- BTEX and styrene (SCC and TCLP)
- TPHs (C10-C36) (SCC and TCLP)

4. Each material was split into 12 kg subsamples for treatment.

5. Subsamples (12 kg) of each material were treated with 5%, 12.5% and 20% cement (by weight). The cement was mixed into the materials by hand, which has previously been demonstrated to replicate a pugmill or equivalent at full-scale treatment.

6. Two cylinders were cast per treatment per material (3 materials x 3 treatments x 2 replicates/treatment = 18 cylinders total).

7. A sufficient quantity of each treated material was separated and stored in tightly sealed plastics bags for five days prior to submitting them for post-treatment chemical analysis.

8. Each cast cylinder was cured and tested in accordance with NSW RTA Test Method T131, in accordance with DECCW General Immobilisation Approval for Coal Tar.

9. One sample from each treatment (total 9) plus 1 untreated control (Material 3) was collected and submitted for chemical analysis. Each sample was analysed for the following:

- As, Cd, Cr, Pb, Hg, Ni (SCC and ASLP)
- Benzo(a)pyrene and PAHs (SCC and ASLP)
- Phenols (SCC and ASLP)
- BTEX and styrene (SCC and ASLP)
- TPHs (C10-C36) (SCC and TCLP)

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10. JBS were unable to provide bulk groundwater samples for the ASLP tests, therefore DI water was used for the ASLP tests.

All chemical analyses were conducted by Envirolab Services and all UCS cylinders were prepared and tested by Douglas Partners Geotechnical Laboratory. Envirolab and Douglas Partners are NATA accredited and conducted the testing in accordance with laboratory testing quality assurance protocols. Laboratory reports are provided in Attachments 1-4.

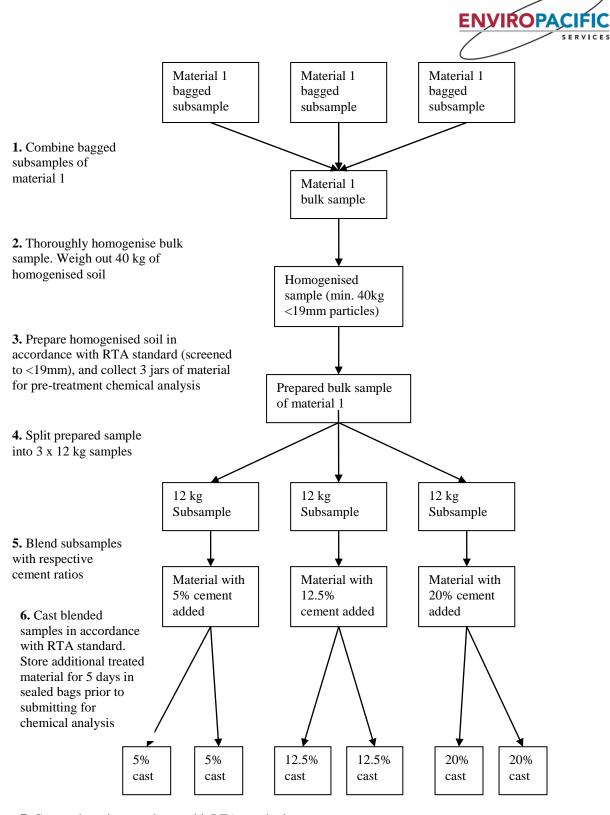
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7. Cure and test in accordance with RTA standard. Undertake chemical analysis on each treated materials following cure period.

Figure 1. Procedural flowchart followed for each material.

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Sample Collection

Bulk sample collection at Macdonaldtown was conducted on 2nd July, 2010 by JBS and subsequently transferred to Enviropacific's Sydney office. Three types of material were targeted:

- Material 1: TP1 at 0.3 0.4 m depth
- Material 2: TP3 at 1.3 1.7 m depth
- Material 3: TP3 at 4.0 4.2 m depth

Material Description

- Material 1 was a silty, gravelly sand fill (refer to Fig 2).
- Material 2 was a silty clay fill (refer to Fig 3).
- Material 3 was a silty clay fill (refer to Fig 4).



Figure 2.

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Figure 3.



Figure 4.

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Results/Discussion

A detailed summary of the chemical analysis and UCS testing results is included in Attachment 5. The total and leachable (TCLP for pre-treated material and ASLP-DI water for post-treated material) concentrations for the main contaminants of concern (BaP, total PAHs and TPH-C10-C36) and UCS results are summarised in Table 1 below. A discussion of the results is provided below.

Table 1. Summary of total and leachable results (TCLP for pre-treated material and ASLP-DI water for post-treated material) for BaP, PAHs and TPH (C10-C36) (ND = Not Determined).

Sample ID	Total BaP (mg/kg)	Leachable BaP TCLP/ ASLP (mg/L)	Total PAHs (mg/kg)	Leachable PAHs TCLP/ ASLP (mg/L)	Total TPH C10-C36 (mg/kg)	Leachable TPH C10-C36 TCLP/ ASLP (µg/L)	Mean UCS 7-day curing (MPa)
General Solid Criteria	10	40	200	-	10,000	-	
Restricted Solid Criteria	23	160	800	-	40,000	-	
Pre-treated materials	•					•	
Material 1 Pre 1	4.4	< 0.001	34	< 0.001	700	120	ND
Material 1 Pre 2	4.5	< 0.001	35	< 0.001	700	110	ND
Material 1 Pre 3	5.6	< 0.001	53	< 0.001	760	<250	ND
Mean	4.8	< 0.001	41	< 0.001	720	115	
Material 2 Pre 1	0.9	< 0.001	276	3.5	650	8740	ND
Material 2 Pre 2	1.1	< 0.001	292	2.8	780	7060	ND
Material 2 Pre 3	1.0	< 0.001	331	2.7	790	6150	ND
Mean	1.0	< 0.001	299	3.0	740	7317	
Material 3 Pre 1	0.2	< 0.001	5	3.3	<250	2120	ND
Material 3 Pre 2	2.1	< 0.001	39	3.0	370	6960	ND
Material 3 Pre 3	1.7	< 0.001	36	1.2	450	1480	ND
Mean	1.3	< 0.001	27	2.5	410	3520	
Post-treated materials	1	1	1	1	-	1	
Control - Material 3	1.7	< 0.001	51	0.083	500	680	ND
Material 1 Post 5%	4.7	< 0.001	44	< 0.001	360	400	2.18
Material 1 Post 12.5%	3.8	< 0.001	34	< 0.001	320	280	3.10
Material 1 Post 20%	3.9	< 0.001	43	< 0.001	300	290	5.85
Material 2 Post 5%	2.0	< 0.001	171	2.6	810	6700	0.35
Material 2 Post 12.5%	1.6	< 0.001	117	1.9	450	5700	1.00
Material 2 Post 20%	1.6	< 0.001	101	1.5	420	4900	1.55
Material 3 Post 5%	0.8	< 0.001	17	0.26	<250	1420	0.13
Material 3 Post 12.5%	0.8	< 0.001	20	0.33	<280	1300	0.43
Material 3 Post 20%	1.7	< 0.001	48	0.48	460	1950	0.60

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Total (SCC) results - untreated samples:

The main contaminants of concern in the materials are total PAHs. The total PAHs in the Material 2 would result in this untreated material being classified as Restricted solid waste, whereas total PAHs in Materials 1 and 3 would result in these untreated materials being classified as General solid waste. Notably, naphthalene comprised ~1%, 90% and 30% of the total PAHs in Materials 1-3, respectively. Generally the percentage of small-ring PAHs in the material reflects both the aging and nature of the material (e.g. the percentage increases as the amount of coal tar material increases). All other contaminants of concern were well below the General solid waste criteria.

Total (SCC) results – cement treated samples:

The total contaminant results for treated samples for each of the materials were generally similar to the untreated sample. Any observed differences appear to be related more to the heterogeneity of the material, rather than dilution of the sample with treatment reagents. The total PAHs in treated samples of Material 2 however decreased by 43% (at 5% cement), 61% (at 12.5% cement) and 66% (at 20% cement), and is mainly attributable to decreased naphthalene in the treated samples of Material 2, which decreased from ~90% of total PAHs to 70% (at 5% cement), 63% (at 12.5% cement) and 59% (at 20% cement). This is most likely related to increased volatilisation of naphthalene with the amount of cement added, resulting from the heat of reaction (i.e. heat of hydration). The percentages of naphthalene in Material 1 and 3 were similar in the treated samples to the untreated samples.

Leachable results – General Comments/Limitations

With the exception of BaP, there are currently no criteria for the leachability of total PAHs or TPHs in either the Waste Classification Guidelines or the General IA for gasworks waste (2005/14). Hence there is no point of reference for the leachability of total PAHs and TPHs provided by the DECCW. The leachable PAHs and TPHs are discussed in this report in the context that this information is considered to be relevant to the retention of treated material on site. With regards to assessing the leachability of contaminants from cement stabilised waste, it is worth noting that current leachate methods (TCLP, ASLP and MEP are the only leachate tests currently used by regulatory authorities in Australia) have important limitations. For example, each of these methods require particle size reduction to either 9.5 mm (TCLP and MEP) or 2.4 mm (ASLP), which effectively contravenes the assessment of encapsulated wastes as the integrity of the monolithic structure is compromised, and each of these methods employ vigorous end-over-end agitation of the sample. In fact in AS4439.3-1997 (ASLP) the Scope states that "The procedure is not applicable to encapsulated wastes which cannot be reduced to the specified maximum particle size without breaking the

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integrity of encapsulation". For this reason, in the General IA, the DECCW is in effect relying on the UCS measurement as an indicator of the stability of cement stabilised waste, whilst still requiring BaP leachability (TCLP) for cement stabilised samples to be below the Waste Classification Guidelines criteria (and from previous EPS experience with other gasworks projects BaP leachability has typically been non-detectable in both the untreated and cement stabilised samples using TCLP).

If a more appropriate leach test (e.g. a diffusion-based or column leach test on a moulded/monolithic sample) was adopted for assessing the leachability of cement stabilised materials destined for on-site placement, the optimum UCS required to minimise contaminant leachability could be more accurately determined, and a different leachability data set may result, that might be more appropriately applied to acceptance criteria for on-site placement.

Leachable results (TCLP) – untreated samples

Leachable results for the large-ring PAHs (>pyrene and including BaP) for all untreated materials were below laboratory detection limits. However, total PAH leachability was relatively high in Materials 2 and 3 (mean results of 3.0 and 2.5 mg/L, respectively) and was mainly attributable to leachable naphthalene (98%). Notably, total naphthalene in these two materials was significantly different (270 mg/kg for Material 2 compared to 6.7 mg/kg for Material 3). The ASLP-DI water total PAH leachability for the untreated control (Material 3) submitted with the treated samples was significantly lower (0.083 mg/L). The leachabilities of all other contaminants of concern for the untreated materials were well below the General solid waste criteria.

Leachable results (ASLP-DI water) – cement treated samples

BaP leachability remained non-detectable in all three materials for all ratios of cement addition. Total PAH leachability remained relatively high in Material 2 (max 2.6 mg/L at 5% cement, min 1.5 mg/L at 20% cement), and was again mainly attributable to leachable naphthalene (~91%). Given the lower total naphthalene in the treated samples for Material 2, there is little evidence of reduced leachability of naphthalene (and other PAHs) in the cement treated samples using the ASLP method, despite achieving UCS strengths of >1 MPa for Material 2 (refer Table 1). In fact, for Material 3 all three cement treated samples demonstrated higher total PAH leachability (up to six times higher for 20% cement) than for the control (untreated Material 3) using ASLP-DI water. Similarly, leachable TPHs for treated samples of Material 3 increased by at least two-fold compared to the untreated control using ASLP-DI water. Increased leachability of organic contaminants from cement stabilised

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samples (and hence lack of evidence of their immobilisation) has been previously observed by EPS in other gasworks projects using TCLP and/or ASLP, and clearly demonstrates the limitations of these methods for assessing the leachability of cement stabilised materials, as mentioned above.

Notably, the leachability of some inorganic contaminants (e.g. arsenic, chromium) were higher in the treated ASLP results compared to untreated TCLP results, which is related to the amphoteric nature of these contaminants where their solubility increases at higher pH. However the leachabilities of all inorganic contaminants in the treated samples were well below the General solid waste criteria.

UCS Results – cement treated samples

UCS testing is currently required by DECCW as part of the General IA for gasworks waste for assessing the stability of cement stabilised wastes destined for off-site landfill disposal with a target of 1 MPa. UCS results for the cement treated materials (refer Table 1) showed that >1 MPa was readily achieved for Material 1 using 5% cement (mean result 2.2 MPa). For Material 2, a higher ratio of cement was required (12.5% cement resulted in 1.0 MPa UCS), which would be related to the higher level of organic contamination in this material and/or the higher clay content. For Material 3, treatment with 20% cement resulted in a UCS of only 0.6 MPa. This sample was observed to have very high clay content, providing very little aggregate for the cement curing process. The low level of organic contaminants observed in Material 3 indicates that the contaminants did not have had a significant effect on the UCS results for this material.

Summary

The level of contamination in the three materials was relatively low for gasworks waste. Total PAHs in the Material 2 would result in the untreated material being classified as Restricted solid waste. All other contaminants of concern were well below the General solid waste criteria. UCS measurements of >1 MPa were achieved for Materials 1 and 2, whereas a maximum UCS of 0.6 MPa was achieved for Material 3.

The results from this trial show that there is no direct relationship between the UCS measurement and the leachability of contaminants from cement stabilised materials. In fact, cement treatment appears to increase the leachability of organic contaminants, as assessed using ASLP-DI water. If a more appropriate (alternative) test was used to assess the leachability of contaminants from stabilised waste to be placed on-site, the optimum UCS required to minimise leachability of contaminants could be more accurately determined.

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Historically in NSW some degree of macroencapsulation of gasworks waste with cement has been undertaken for waste being disposed of to landfill, in order to provide a sufficient level of confidence in the long-term stability of the treated waste.

Attachments

Attachments 1-3 - Laboratory reports from Envirolab Services

Attachment 4 – Laboratory reports from Douglas Partners

Attachment 5 – Macdonaldtown Results Summary

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DETERMINATION OF UNCONFINED COMPRESSIVE STRENGTH OF **COMPACTED MATERIALS**

	Client: Enviropacific Services	Enviropacific Services Pty Ltd		66360 S10-006 D	
Îă −ur urați	Project: Macdonaldtown Gaswork	<s< th=""><th>Report No: Report Date:</th><th>04/08/10</th></s<>	Report No: Report Date:	04/08/10	
· ·	Location: Macdonaldtown		Date Sampled: Date of Test: Page:	27/07/10 27/07/10 1 of 1	
	Material Retained on 19mm sieve:	0%			
	Material Description:	Material 1 + 5% cem	nent		
-					
	Elapsed time between addition of binder and compaction:	15 mins			
	Method of Compaction: No. of layers	Standard 3			
	Curing Details:	7 day accelerated cu	rated curing at 65°C & +-5°C		
	:	Specimen A	Specime	n B	
	Moisture Content at Compaction:	14.5%	14.0%		
1 .	Load at failure kN:	16.576%	21.042%		
	Dry Density of test specimens:	1.83t/m³	1.86t/m³		
	Unconfined Compressive Strength:	1.90MPa	2.45MPa		

Test Method(s): RTA T116, T131 Sampled by client - compacted in laboratory Sampling Method(s): **Remarks:**

Approved Signatory:

Tested: MBG

Checked. NW

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Norman Weimann Laboratory Manager



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DETERMINATION OF UNCONFINED COMPRESSIVE STRENGTH OF **COMPACTED MATERIALS**

Client:	Enviropacific Services Pty	Project No: Report No:	66360 S10-006 E	
Project:	Macdonaldtown Gasworks		Report Date:	04/08/10
Location:	Macdonaldtown		Date Sampled: Date of Test: Page:	27/07/10 27/07/10 1 of 1
Material Retai	ned on 19mm sieve:	0%		
Material Desc	ription:	Material 1 + 12.5% cei	ment	
Elapsed time binder and co	between addition of mpaction:	15 mins		
Method of Compaction: No. of layers		Standard 3		
Curing Details	5:	7 day accelerated curing at 65°C & +-5°C		
		Specimen C	Specime	n D
Moisture Cont	tent at Compaction:	12.5%	12.5%	
Load at failure	e kN:	28.914%	25.484%	
Dry Density o	f test specimens:	1.86t/m³	1.84t/m³	
Unconfined C	ompressive Strength:	3.30MPa	2.90MPa	

RTA T116, T131 Test Method(s): Sampled by client - compacted in laboratory Sampling Method(s): **Remarks:**

Approved Signatory:

Tested: MBG

Checked: NW

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Norman Weimann Laboratory Manager



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DETERMINATION OF UNCONFINED COMPRESSIVE STRENGTH OF COMPACTED MATERIALS

Client:	Enviropacific Services Pty Ltd		Project No:	66360
Project:	Macdonaldtown Gasworks		Report No: Report Date:	S10-006 F 04/08/10
Location:	Macdonaldtown		Date Sampled: Date of Test: Page:	27/07/10 27/07/10 1 of 1
Material Retai	ned on 19mm sieve:	0%		
Material Desci	iption:	Material 1 + 20.0% cer	nent	
Elapsed time between addition of binder and compaction:		15 mins Standard		
Method of Cor No. of layers		3		
Curing Details	:	7 day accelerated curing at 65°C & +-5°C		
		Specimen E	Specimer	ı F
Moisture Conte	ent at Compaction:	11.5%	11.5%	
Load at failure	kN:	53.145%	47.990%	
Dry Density of	test specimens:	1.82t/m³	1.85t/m³	
Unconfined Co	ompressive Strength:	6.15MPa	5.55MPa	
Moisture Content at Compaction: Load at failure kN: Dry Density of test specimens: Unconfined Compressive Strength:		11.5% 53.145% 1.82t/m³	11.5% 47.990% 1.85t/m³	۶F

 Test Method(s):
 RTA T116, T131

 Sampling Method(s):
 Sampled by client – compacted in laboratory

 Remarks:
 Remarks:

Approved Signatory:

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Testeo:

Checked: NW

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Norman Weimann Laboratory Manager



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 (02) 9809 0666

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 sydney@douglespartners.com.au

DETERMINATION OF UNCONFINED COMPRESSIVE STRENGTH OF COMPACTED MATERIALS

Client: Enviropacific Services Pty		y Ltđ	Project No:	66360
Project:	Macdonaldtown Gasworks		Report No: Report Date:	S10-006 G 06/08/10
Location:	Macdonaldtown	· · · · ·	Date Sampled: Date of Test: Page:	30/07/10 .30/07/10 1 of 1
Material Retair	ned on 19mm sieve:	0%		i the defaure of the second
Material Descr	iption:	Material 2 + 5%		
Elapsed time b binder and con	petween addition of npaction:	15 mins		
Method of Con No. of layers	npaction:	Standard 3		
Curing Details:		7 day accelerated curin	ng at 65°C & +-5°C	c
		Specimen A	Specimen	В
Moisture Conte	ent at Compaction:	16.5%	17.0%	
Load at failure	kN:	3.008%	3.018%	
Dry Density of	test specimens:	1.76t/m³	1.74t/m³	
Unconfined Co	mpressive Strength:	0.35MPa	0.35MPa	
	······································			

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 Test Method(s):
 RTA T116, T131

 Sampling Method(s):
 Sampled by client – compacted in laboratory

 Remarks:
 Remarks:

Approved Signatory:

Tested: MBG

Checked: NW

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Norman Weimann Laboratory Manager



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DETERMINATION OF UNCONFINED COMPRESSIVE STRENGTH OF COMPACTED MATERIALS

Client:	Enviropacific Services Pty Ltd		Project No:	66360
Project:	Macdonaldtown Gasworks		Report No: Report Date:	S10-006 H 06/08/10
Location:	Macdonaldtown		Date Sampled: Date of Test: Page:	30/07/10 30/07/10 1 of 1
Material Retair	ned on 19mm sieve:	0%		,
Material Descr	iption:	Material 2 + 12.5%		
Elapsed time between addition of binder and compaction: Method of Compaction:		15 mins Standard		
No. of layers Curing Details:		3		_
Curing Details.		7 day accelerated curir	-	
Moisture Conte	ent at Compaction:	Specimen C	Specimen	D
Load at failure	•	15.5%	15.0%	
		8.928%	8.353%	
Dry Density of	test specimens:	1.77t/m³	1.72t/m³	
Unconfined Co	mpressive Strength:	1.05MPa	0.95MPa	

Test Method(s):RTA T146, T131Sampling Method(s):Sampled by client – compacted in laboratoryRemarks:

Approved Signatory:

MBG

Tested:

Checked: NW

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Norman Weimann Laboratory Manager



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DETERMINATION OF UNCONFINED COMPRESSIVE STRENGTH OF **COMPACTED MATERIALS**

Client: Enviropacific Services I		y Ltd	Project No:	66360
Project:	Macdonaldtown Gasworks		Report No: Report Date:	S10-006 I 06/08/10
Location:	Macdonaldtown		Date Sampled: Date of Test: Page:	30/07/10 30/07/10 1 of 1
Material Retain	ned on 19mm sieve:	0%		
Material Descr	iption:	Material 2 + 20.0%		
Elapsed time between addition of binder and compaction: Method of Compaction: No. of layers		15 mins Standard 3		
Curing Details:		7 day accelerated curing at 65°C & +-5°C		
		Specimen E	Specimen	F
Moisture Conte	ent at Compaction:	14.5%	13.5%	
Load at failure	kN:	14.184%	12.705%	
Dry Density of	test specimens:	1.76t/m³	1.76t/m³	
Unconfined Co	mpressive Strength:	1.65MPa	1.45MPa	
			· · · · · · · · · · · · · · · · · · ·	-

Test Method(s): RTA T116, T131 Sampling Method(s): Sampled by client - compacted in laboratory **Remarks:**

Approved Signatory:

M8G

Tested:

Checked: NW

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Norman Weimann Laboratory Manager



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DETERMINATION OF UNCONFINED COMPRESSIVE STRENGTH OF COMPACTED MATERIALS

Client:	Enviropacific Services Pty Ltd		Project No:	66360 S10-006 A
Project:	Macdonaldtown Gasworks		Report No: Report Date:	04/08/10
Location:	Macdonaldtown		Date Sampled: Date of Test: Page:	27/07/10 27/07/10 1 of 1
Material Retai	ned on 19mm sieve:	0%		
Material Desc	ription:	Material 3 + 5% ceme	nt	
Elapsed time binder and co	between addition of mpaction:	15 mins		
Method of Col No. of layers	mpaction:	Standard 3		
Curing Details	::	7 day accelerated curing at 65°C & +-5°C		
		Specimen A	Specimer	ηB
Moisture Cont	ent at Compaction:	27.5%	26.5%	
Load at failure	kN:	0.894%	1.417%	
Dry Density of	f test specimens:	1.57t/m³	1.58t/m³	
Unconfined C	ompressive Strength:	0.10MPa	0.15MPa	

Test Method(s):RTA T116, T131Sampling Method(s):Sampled by client – compacted in laboratoryRemarks:

Approved Signatory:

MBG

Tested:

Checked, NW

Magnares Norman Weimann

Laboratory Manager

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DETERMINATION OF UNCONFINED COMPRESSIVE STRENGTH OF **COMPACTED MATERIALS**

Client: Enviropacific Services Pty		y Ltd	Project No:	66360	
Project:	Macdonaldtown Gasworks		Report No: Report Date:	S10-006 B 04/08/10	
Location:	Macdonaldtown		Date Sampled: Date of Test: Page:	27/07/10 27/07/10 1 of 1	
Material Retail	ned on 19mm sieve:	0%			
Material Descr	ription:	Material 3 + 12.5% cer	nent		
Elapsed time between addition of binder and compaction:		15 mins			
Method of Compaction: No. of layers		Standard 3			
Curing Details	:	7 day accelerated curing at 65°C & +- 5°C			
		Specimen C	Specimen	۱D	
Moisture Conte	ent at Compaction:	26.0%	26.0%		
Load at failure	kN:	3.991%	3.395%		
Dry Density of	test specimens:	1.60t/m³	1.59t/m³		
Unconfined Co	ompressive Strength:	0.45MPa	0.40MPa		

Test Method(s): RTA T116, T131 Sampling Method(s): Sampled by client - compacted in laboratory **Remarks:**

Approved Signatory:

Tested: MBG Checked: NW Millermany Norman Weimann Laboratory Manager



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DETERMINATION OF UNCONFINED COMPRESSIVE STRENGTH OF COMPACTED MATERIALS

Client: Enviropacific Services		Pty Ltd	Project No:	66360
Project:	Macdonaldtown Gasworks	6	Report No: Report Date:	S10-006 C 04/08/10
Location:	Macdonaldtown	· · · ·	Date Sampled: Date of Test: Page:	27/07/10 27/07/10 1 of 1
Material Retai	ned on 19mm sieve:	0%		
Material Desci	ription:	Material 3 + 20% ceme	ent	
Elapsed time t binder and cor	between addition of npaction:	15 mins		
Method of Con No. of layers	npaction:	Standard 3		
Curing Details:		7 day accelerated curin	ng at 65°C & +-5°C	>
		Specimen E	Specimen	F
Moisture Conte	ent at Compaction:	22.0%	24.0%	
Load at failure	kN:	4.879%	5.588%	
Dry Density of	test specimens:	1.62t/m³	1.60t/m ³	
Unconfined Co	mpressive Strength:	0.55MPa	0.65MPa	

Test Method(s):RTA T116, T131Sampling Method(s):Sampled by client – compacted in laboratoryRemarks:

Approved Signatory:

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Checked: NW

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Norman Weimann Laboratory Manager



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Envirolab Services Pty Ltd ABN 37 112 535 645 12 Ashley St Chatswood NSW 2067 ph 02 9910 6200 fax 02 9910 6201 enquiries@envirolabservices.com.au www.envirolabservices.com.au

CERTIFICATE OF ANALYSIS 43953

Client: Enviropacific Services (Chatswood) Pty Ltd 1/28 Barcoo St Chatswood NSW 2067

Attention: Marty Croker

Sample log in details:

Your Reference: No. of samples: Date samples received: Date completed instructions received:

E10100 Macdonaldtown Gasworks 9 Soils 26/07/10 26/07/10

Analysis Details:

Please refer to the following pages for results, methodology summary and quality control data. Samples were analysed as received from the client. Results relate specifically to the samples as received. Results are reported on a dry weight basis for solids and on an as received basis for other matrices. *Please refer to the last page of this report for any comments relating to the results.*

Report Details:

 Date results requested by:
 2/08/10

 Date of Preliminary Report:
 Not Issued

 Issue Date:
 13/08/10

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 Tests not covered by NATA are denoted with *.

Results Approved By:

Giovanni Agosti Technical Manager

Sandra Tastor Senior Organic Chemist

Jacinta Hurst Laboratory Manager

Envirolab Reference: 43953 Revision No: R 01

Nancy Zhang Chemist



Page 1 of 30

Client Reference: E10100 Macdonaldtown Gasworks

VOCs in soil						
Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pre
		1	2	3	1	2
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	31/7/2010	31/7/2010	31/7/2010	31/7/2010	31/7/2010
styrene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Surrogate Dibromofluorometha	%	93	91	90	92	92
Surrogate aaa-Trifluorotoluene	%	70	77	75	75	72
Surrogate Toluene-da	%	110	107	96	112	113
Surrogate 4-Bromofluorobenzene	%	94	93	92	110	103

VOCs in soil					
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9
Your Reference		Material 2 Pre	Material 3 Pre	Material 3 Pre	Material 3 Pre
		3	1	2	3
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	31/7/2010	31/7/2010	31/7/2010	31/7/2010
styrene	mg/kg	<1.0	<1.0	<1.0	<1.0
Surrogate Dibromofluorometha	%	95	95	96	95
Surrogate aaa-Trifluorotoluene	%	73	73	73	72
Surrogate Toluene-da	%	114	114	114	113
Surrogate 4-Bromofluorobenzene	%	105	96	95	96

ACCREDITED FOR TECHNICAL COMPETENCE

Client Reference: E10100 Macdonaldtown Gasworks

vTPH & BTEX in Soil						
Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pr
		1	2	3	1	2
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	30/7/2010	30/7/2010	30/7/2010	30/7/2010	30/7/2010
vTPH C6 - C9	mg/kg	<25	<25	<25	54	37
Benzene	mg/kg	<0.5	<0.5	<0.5	<0.5	<0.5
Toluene	mg/kg	<0.5	<0.5	<0.5	0.70	0.60
Ethylbenzene	mg/kg	<1.0	<1.0	<1.0	19	14
m+p-xylene	mg/kg	<2.0	<2.0	<2.0	16	11
o-Xylene	mg/kg	<1.0	<1.0	<1.0	15	11
Surrogate aaa-Trifluorotoluene	%	70	77	75	75	72

vTPH & BTEX in Soil					
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9
Your Reference		Material 2 Pre	Material 3 Pre	Material 3 Pre	Material 3 Pre
		3	1	2	3
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	30/7/2010	30/7/2010	30/7/2010	30/7/2010
vTPH C6 - C9	mg/kg	65	<25	<25	<25
Benzene	mg/kg	<0.5	<0.5	<0.5	<0.5
Toluene	mg/kg	0.80	<0.5	<0.5	<0.5
Ethylbenzene	mg/kg	21	<1.0	<1.0	<1.0
m+p-xylene	mg/kg	17	<2.0	<2.0	<2.0
o-Xylene	mg/kg	15	<1.0	<1.0	<1.0
Surrogate aaa-Trifluorotoluene	%	73	73	73	72

R 01



Client Reference: E10100 Macdonaldtown Gasworks

sTPH in Soil (C10-C36)						
Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pre
		1	2	3	1	2
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	29/7/2010
TPH C10 - C14	mg/kg	<50	<50	<50	440	580
TPH C15 - C28	mg/kg	320	330	390	210	200
TPH C29 - C36	mg/kg	380	370	370	<100	<100
Surrogate o-Terphenyl	%	#	#	#	#	#
sTPH in Soil (C10-C36)						
Our Reference	LINITS	43953-6	43953-7	43953-8	43953-9	

STPH IN SOIL (CTU-C36)					
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9
Your Reference		Material 2 Pre	Material 3 Pre	Material 3 Pre	Material 3 Pre
		3	1	2	3
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010
TPH C10 - C14	mg/kg	570	<50	110	140
TPH C15 - C28	mg/kg	220	<100	260	310
TPH C29 - C36	mg/kg	<100	<100	<100	<100
Surrogate o-Terphenyl	%	#	100	#	#

Envirolab Reference: 43953 **Revision No:**

R 01



	Clie	ent Reference	e: E10100				
PAHs in Soil							
Our Reference:		UNITS	43953-1	43953-2	43953-3	43953-4	
Vaux Deference			Motorial 1 Dra	Motorial 1 Dra	Matarial 1 Dra	Matarial 2 Dra	140

Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pre
		1	2	3	1	2
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Date analysed	-	30/07/2010	30/07/2010	30/07/2010	30/07/2010	30/07/2010
Naphthalene	mg/kg	0.4	0.4	0.5	250	260
Acenaphthylene	mg/kg	0.7	0.7	1.4	1.6	2.0
Acenaphthene	mg/kg	<0.1	<0.1	<0.1	3.7	4.9
Fluorene	mg/kg	0.2	0.2	0.5	3.3	4.2
Phenanthrene	mg/kg	2.0	1.8	5.8	5.8	7.1
Anthracene	mg/kg	0.6	0.6	1.5	1.6	2.0
Fluoranthene	mg/kg	4.3	4.7	8.3	2.4	2.8
Pyrene	mg/kg	5.0	6.2	8.5	3.0	3.4
Benzo(a)anthracene	mg/kg	3.3	3.3	4.4	1	1.1
Chrysene	mg/kg	2.7	3.2	4.0	0.9	1.1
Benzo(b+k)fluoranthene	mg/kg	6.0	5.5	7.5	1.0	1.2
Benzo(a)pyrene	mg/kg	4.4	4.5	5.6	0.9	1.1
Indeno(1,2,3-c,d)pyrene	mg/kg	2.0	1.9	2.4	0.3	0.3
Dibenzo(a,h)anthracene	mg/kg	0.5	0.4	0.5	<0.1	<0.1
Benzo(g,h,i)perylene	mg/kg	1.9	2.0	2.3	0.3	0.4
Surrogate p-Terphenyl-d14	%	75	77	74	80	66



PAHs in Soil					
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9
Your Reference		Material 2 Pre	Material 3 Pre	Material 3 Pre	Material 3 Pre
		3	1	2	3
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil
Date extracted	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Date analysed	-	30/07/2010	30/07/2010	30/07/2010	30/07/2010
Naphthalene	mg/kg	300	1.7	11	7.4
Acenaphthylene	mg/kg	1.9	0.1	0.8	0.9
Acenaphthene	mg/kg	4.6	0.1	0.4	0.7
Fluorene	mg/kg	4.0	0.2	1.3	1.7
Phenanthrene	mg/kg	6.9	0.6	4.3	5.2
Anthracene	mg/kg	1.9	0.2	1.2	1.5
Fluoranthene	mg/kg	2.8	0.4	4.1	4.1
Pyrene	mg/kg	3.5	0.6	5.0	5.3
Benzo(a)anthracene	mg/kg	1.1	0.2	2.2	1.9
Chrysene	mg/kg	1.1	0.2	2.1	1.8
Benzo(b+k)fluoranthene	mg/kg	1.1	0.3	2.9	2.3
Benzo(a)pyrene	mg/kg	1.0	0.2	2.1	1.7
Indeno(1,2,3-c,d)pyrene	mg/kg	0.3	<0.1	0.8	0.6
Dibenzo(a,h)anthracene	mg/kg	<0.1	<0.1	0.2	0.2
Benzo(g,h,i)perylene	mg/kg	0.3	<0.1	0.9	0.7
Surrogate p-Terphenyl-d14	%	76	75	79	92

Envirolab Reference: 43953 Revision No: R 01

Client Reference: E10100 Macdonaldtown Gasworks	Client Reference:	E10100 Macdonaldtown Gasworks
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Speciated Phenols in Soil						
Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pre
		1	2	3	1	2
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	2/8/2010	2/8/2010	2/8/2010	2/8/2010	2/8/2010
Phenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2-Chlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2-Methylphenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
3/4-Methylphenol	mg/kg	<2.0	<2.0	<2.0	<2.0	<2.0
2-Nitrophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,4-Dimethylphenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,4-Dichlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,6-dichlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,4,5-trichlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,4,6-trichlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,4-dinitrophenol	mg/kg	<10	<10	<10	<10	<10
4-nitrophenol	mg/kg	<10	<10	<10	<10	<10
2,3,4,6-tetrachlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2-methyl-4,6-dinitrophenol	mg/kg	<10	<10	<10	<10	<10
pentachlorophenol	mg/kg	<10	<10	<10	<10	<10
Surrogate 2-fluorophenol	%	105	95	92	116	123
Surrogate Phenol-d6	%	68	84	82	96	89
Surrogate 2,4,6-Tribromophenol	%	93	99	89	139	140
Surrogate p-Terphenyl-d14	%	114	114	105	116	114

Client Reference:	E10100 Macdonaldtown Gasworks

Speciated Phenols in Soil					
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9
Your Reference		Material 2 Pre	Material 3 Pre	Material 3 Pre	Material 3 Pre
Date Sampled		3 26/07/2010	1 26/07/2010	2 26/07/2010	3 26/07/2010
Type of sample		Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	2/8/2010	2/8/2010	2/8/2010	2/8/2010
Phenol	mg/kg	<1.0	<1.0	<1.0	<1.0
2-Chlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0
2-Methylphenol	mg/kg	<1.0	<1.0	<1.0	<1.0
3/4-Methylphenol	mg/kg	<2.0	<2.0	<2.0	<2.0
2-Nitrophenol	mg/kg	<1.0	<1.0	<1.0	<1.0
2,4-Dimethylphenol	mg/kg	<1.0	<1.0	<1.0	<1.0
2,4-Dichlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0
2,6-dichlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0
2,4,5-trichlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0
2,4,6-trichlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0
2,4-dinitrophenol	mg/kg	<10	<10	<10	<10
4-nitrophenol	mg/kg	<10	<10	<10	<10
2,3,4,6-tetrachlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0
2-methyl-4,6-dinitrophenol	mg/kg	<10	<10	<10	<10
pentachlorophenol	mg/kg	<10	<10	<10	<10
Surrogate 2-fluorophenol	%	122	111	116	72
Surrogate Phenol-de	%	82	72	122	63
Surrogate 2,4,6-Tribromophenol	%	127	100	120	78
Surrogate p-Terphenyl-d14	%	108	106	110	76

Acid Extractable metals in soil						
Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pi
		1	2	3	1	2
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date digested	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Date analysed	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Arsenic	mg/kg	36	38	44	<4	<4
Cadmium	mg/kg	1.1	1.4	1.3	<0.5	<0.5
Chromium	mg/kg	19	20	20	14	14
Lead	mg/kg	230	230	300	27	32
Mercury	mg/kg	0.4	0.3	0.4	<0.1	<0.1
Nickel	mg/kg	21	26	27	3	3
Nickel Acid Extractable metals in soil	mg/kg	21	26	27	3	
		10050.0	10050 5	10050.0	10050.0	

Acid Extractable metals in soil					
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9
Your Reference		Material 2 Pre	Material 3 Pre	Material 3 Pre	Material 3 Pre
		3	1	2	3
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil
Date digested	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Date analysed	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Arsenic	mg/kg	8	<4	5	6
Cadmium	mg/kg	<0.5	<0.5	<0.5	<0.5
Chromium	mg/kg	19	14	20	24
Lead	mg/kg	43	11	21	29
Mercury	mg/kg	<0.1	<0.1	<0.1	<0.1
Nickel	mg/kg	3	2	2	3



Moisture						
Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pre
		1	2	3	1	2
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date prepared	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Moisture	%	15	10	16	21	21
	1	T	T	T	1	
Moisture						
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9	
Your Reference		Material 2 Pre	Material 3 Pre	Material 3 Pre	Material 3 Pre	
		3	1	2	3	
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	
Type of sample		Soil	Soil	Soil	Soil	
Date prepared	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	
Date analysed	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	
Moisture	%	2.7	21	22	20	

Envirolab Reference: 43953 **Revision No:**



VOCs in Zero Headspace						
Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pre
		1	2	3	1	2
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	31/7/2010	31/7/2010	31/7/2010	31/7/2010	31/7/2010
Styrene	µg/L	<1.0	<1.0	<1.0	3.2	3.0
Surrogate Dibromofluoromethane	%	107	109	111	108	104
Surrogate toluene-d8	%	61	110	97	112	99
Surrogate 4-BFB	%	95	93	94	121	115

VOCs in Zero Headspace					
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9
Your Reference		Material 2 Pre 3	Material 3 Pre 1	Material 3 Pre 2	Material 3 Pre 3
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	31/7/2010	31/7/2010	31/7/2010	31/7/2010
Styrene	µg/L	2.3	1.2	1.3	1.2
Surrogate Dibromofluoromethane	%	100	98	106	107
Surrogate toluene-d8	%	102	117	108	100
Surrogate 4-BFB	%	114	103	105	107

Envirolab Reference: 43953 **Revision No:**



BTEX in Zero Headspace						
Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pre
		1	2	3	1	2
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	31/7/2010	31/7/2010	31/7/2010	31/7/2010	31/7/2010
Benzene	µg/L	<1.0	<1.0	<1.0	10	10
Toluene	µg/L	<1.0	<1.0	<1.0	19	17
Ethylbenzene	µg/L	<1.0	<1.0	<1.0	270	260
m+p-xylene	µg/L	<2.0	<2.0	<2.0	250	240
o-xylene	µg/L	<1.0	<1.0	<1.0	210	200
Surrogate Dibromofluoromethane	%	107	109	111	108	104
Surrogate toluene-d8	%	101	110	97	112	99
Surrogate 4-BFB	%	95	93	94	121	115

BTEX in Zero Headspace					
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9
Your Reference		Material 2 Pre	Material 3 Pre	Material 3 Pre	Material 3 Pre
		3	1	2	3
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	31/7/2010	31/7/2010	31/7/2010	31/7/2010
Benzene	µg/L	13	<1.0	<1.0	<1.0
Toluene	µg/L	21	<1.0	<1.0	<1.0
Ethylbenzene	µg/L	290	3.7	8.1	8.8
m+p-xylene	µg/L	270	2.2	5.0	6.1
o-xylene	µg/L	230	2.7	5.9	6.8
Surrogate Dibromofluoromethane	%	100	98	106	107
Surrogate toluene-d8	%	102	117	108	100
Surrogate 4-BFB	%	114	103	105	107

Metals in TCLP USEPA1311						
Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pre
		1	2	3	1	2
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Date analysed	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010	29/07/2010
pH of soil for fluid# determ.	pH units	6.80	7.40	7.70	7.90	7.60
pH of soil for fluid # determ. (acid)	pH units	1.50	1.50	1.50	1.50	1.40
Extraction fluid used	-	1	1	1	1	1
pH of final Leachate	pH units	5.10	5.10	5.10	5.00	5.00
Arsenic in TCLP	mg/L	<0.05	<0.05	<0.05	<0.05	<0.05
Cadmium in TCLP	mg/L	<0.01	<0.01	<0.01	<0.01	<0.01
Chromium in TCLP	mg/L	<0.01	<0.01	<0.01	<0.01	<0.01
Lead in TCLP	mg/L	0.04	0.07	0.03	<0.03	<0.03
Mercury in TCLP	mg/L	<0.0005	<0.0005	<0.0005	<0.0005	<0.0005
Nickel in TCLP	mg/L	<0.02	<0.02	<0.02	<0.02	<0.02

Metals in TCLP USEPA1311					
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9
Your Reference		Material 2 Pre	Material 3 Pre	Material 3 Pre	Material 3 Pre
		3	1	2	3
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil
Date extracted	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Date analysed	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010
pH of soil for fluid# determ.	pH units	7.40	5.00	6.10	5.90
pH of soil for fluid # determ. (acid)	pH units	1.50	1.40	1.40	1.40
Extraction fluid used	-	1	1	1	1
pH of final Leachate	pH units	5.00	4.90	4.90	4.90
Arsenic in TCLP	mg/L	<0.05	<0.05	<0.05	<0.05
Cadmium in TCLP	mg/L	<0.01	<0.01	<0.01	<0.01
Chromium in TCLP	mg/L	<0.01	<0.01	<0.01	<0.01
Lead in TCLP	mg/L	<0.03	<0.03	<0.03	0.05
Mercury in TCLP	mg/L	<0.0005	<0.0005	<0.0005	<0.0005
Nickel in TCLP	mg/L	<0.02	<0.02	<0.02	<0.02

Envirolab Reference: 43953 **Revision No:**



TPH in TCLP extract						
Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pre
		1	2	3	1	2
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010	29/7/2010
TPH C10 - C14	µg/L	<50	<50	<50	8,000	6,500
TPH C15 - C28	µg/L	120	110	<100	740	560
TPH C29 - C36	µg/L	<100	<100	<100	<100	<100
Surrogate o-Terphenyl	%	120	129	109	#	#

TPH in TCLP extract					
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9
Your Reference		Material 2 Pre	Material 3 Pre	Material 3 Pre	Material 3 Pre
		3	1	2	3
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil
Date extracted	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010
Date analysed	-	29/7/2010	29/7/2010	29/7/2010	29/7/2010
TPH C10 - C14	µg/L	5,700	1,700	6,000	990
TPH C15 - C28	µg/L	450	420	960	490
TPH C ₂₉ - C ₃₆	µg/L	<100	<100	<100	<100
Surrogate o-Terphenyl	%	#	#	#	123



PAHs in TCLP (USEPA 1311)						
Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pre
Data Oriented		1	2	3	1	2
Date Sampled		26/07/2010 Soil	26/07/2010 Soil	26/07/2010 Soil	26/07/2010 Soil	26/07/2010 Soil
Type of sample		5011	5011	501	501	5011
Date extracted	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Date analysed	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Naphthalene in TCLP	mg/L	<0.001	<0.001	<0.001	3.4	2.7
Acenaphthylene in TCLP	mg/L	<0.001	<0.001	<0.001	0.03	0.02
Acenaphthene in TCLP	mg/L	<0.001	<0.001	<0.001	0.040	0.030
Fluorene in TCLP	mg/L	<0.001	<0.001	<0.001	0.020	0.020
Phenanthrene in TCLP	mg/L	<0.001	<0.001	<0.001	0.010	0.010
Anthracene in TCLP	mg/L	<0.001	<0.001	<0.001	0.002	0.002
Fluoranthene in TCLP	mg/L	<0.001	<0.001	<0.001	0.001	<0.001
Pyrene in TCLP	mg/L	<0.001	<0.001	<0.001	0.001	<0.001
Benzo(a)anthracene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Chrysene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Benzo(b+k)fluoranthene in TCLP	mg/L	<0.002	<0.002	<0.002	<0.002	<0.002
Benzo(a)pyrene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Indeno(1,2,3-c,d)pyrene - TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Dibenzo(a,h)anthracene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Benzo(g,h,i)perylene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Surrogate p-Terphenyl-d14	%	103	108	93	85	129



	Client Reference:	E10100 Macdonaldtown Gasworks
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PAHs in TCLP (USEPA 1311)					
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9
Your Reference		Material 2 Pre	Material 3 Pre	Material 3 Pre	Material 3 Pre
		3	1	2	3
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil
Date extracted	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Date analysed	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Naphthalene in TCLP	mg/L	2.6	3.2	2.9	1.2
Acenaphthylene in TCLP	mg/L	0.02	0.001	0.003	0.001
Acenaphthene in TCLP	mg/L	0.030	0.009	0.010	0.007
Fluorene in TCLP	mg/L	0.020	0.020	0.020	0.010
Phenanthrene in TCLP	mg/L	0.010	0.020	0.020	0.010
Anthracene in TCLP	mg/L	0.002	0.003	0.003	0.002
Fluoranthene in TCLP	mg/L	0.001	0.002	0.002	0.002
Pyrene in TCLP	mg/L	0.001	0.002	0.002	0.002
Benzo(a)anthracene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001
Chrysene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001
Benzo(b+k)fluoranthene in TCLP	mg/L	<0.002	<0.002	<0.002	<0.002
Benzo(a)pyrene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001
Indeno(1,2,3-c,d)pyrene - TCLP	mg/L	<0.001	<0.001	<0.001	<0.001
Dibenzo(a,h)anthracene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001
Benzo(g,h,i)perylene in TCLP	mg/L	<0.001	<0.001	<0.001	<0.001
Surrogate p-Terphenyl-d14	%	90	104	86	101



Client Reference: E10100 Macdonaldtown Gaswork
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Speciated Phenols in water						
Our Reference:	UNITS	43953-1	43953-2	43953-3	43953-4	43953-5
Your Reference		Material 1 Pre	Material 1 Pre	Material 1 Pre	Material 2 Pre	Material 2 Pre
		1	2	3	1	2
Date Sampled		26/07/2010	26/07/2010	26/07/2010	26/07/2010	26/07/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Date analysed	-	30/07/2010	30/07/2010	30/07/2010	30/07/2010	30/07/2010
Phenol	µg/L	<10	<10	<10	<10	<10
2-Chlorophenol	µg/L	<10	<10	<10	<10	<10
2-Methylphenol	µg/L	<10	<10	<10	<10	<10
3/4-Methylphenol	µg/L	<20	<20	<20	<20	<20
2-Nitrophenol	µg/L	<10	<10	<10	<10	<10
2,4-Dimethylphenol	µg/L	<10	<10	<10	<10	<10
2,4-Dichlorophenol	µg/L	<10	<10	<10	<10	<10
2,6-Dichlorophenol	µg/L	<10	<10	<10	<10	<10
2,4,5-Trichlorophenol	µg/L	<10	<10	<10	<10	<10
2,4,6-Trichlorophenol	µg/L	<10	<10	<10	<10	<10
2,4-Dinitrophenol	µg/L	<100	<100	<100	<100	<100
4-Nitrophenol	µg/L	<100	<100	<100	<100	<100
2,3,4,6-Tetrachlorophenol	µg/L	<10	<10	<10	<10	<10
2-methyl-4,6-dinitrophenol	µg/L	<100	<100	<100	<100	<100
Pentachlorophenol	µg/L	<100	<100	<100	<100	<100
Surrogate 2-fluorophenol	%	68	59	54	90	66
Surrogate Phenol-d6	%	52	44	42	49	35
Surrogate 2,4,6-Tribromophenol	%	131	128	105	116	101
Surrogate p-Terphenyl-d14	%	85	93	95	75	60

Client Reference:	E10100 Macdonaldtown Gasworks

Speciated Phenols in water					
Our Reference:	UNITS	43953-6	43953-7	43953-8	43953-9
Your Reference		Material 2 Pre	Material 3 Pre	Material 3 Pre	Material 3 Pre
		3	1	2	3
Date Sampled Type of sample		26/07/2010 Soil	26/07/2010 Soil	26/07/2010 Soil	26/07/2010 Soil
Date extracted	-	29/07/2010	29/07/2010	29/07/2010	29/07/2010
Date analysed	-	30/07/2010	30/07/2010	30/07/2010	30/07/2010
Phenol	µg/L	<10	<10	<10	<10
2-Chlorophenol	µg/L	<10	<10	<10	<10
2-Methylphenol	µg/L	<10	<10	<10	<10
3/4-Methylphenol	µg/L	<20	<20	<20	<20
2-Nitrophenol	µg/L	<10	<10	<10	<10
2,4-Dimethylphenol	µg/L	<10	<10	<10	<10
2,4-Dichlorophenol	µg/L	<10	<10	<10	<10
2,6-Dichlorophenol	µg/L	<10	<10	<10	<10
2,4,5-Trichlorophenol	µg/L	<10	<10	<10	<10
2,4,6-Trichlorophenol	µg/L	<10	<10	<10	<10
2,4-Dinitrophenol	µg/L	<100	<100	<100	<100
4-Nitrophenol	µg/L	<100	<100	<100	<100
2,3,4,6-Tetrachlorophenol	µg/L	<10	<10	<10	<10
2-methyl-4,6-dinitrophenol	µg/L	<100	<100	<100	<100
Pentachlorophenol	µg/L	<100	<100	<100	<100
Surrogate 2-fluorophenol	%	59	59	66	58
Surrogate Phenol-d6	%	32	46	45	46
Surrogate 2,4,6-Tribromophenol	%	71	126	121	121
Surrogate p-Terphenyl-d14	%	89	91	84	86

Method ID	Methodology Summary
GC.14	Soil samples are extracted with methanol and spiked into water prior to analysing by purge and trap GC-MS.
GC.16	Soil samples are extracted with methanol and spiked into water prior to analysing by purge and trap GC-MS. Water samples are analysed directly by purge and trap GC-MS.
GC.3	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-FID.
GC.12 subset	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-MS.
GC.12	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-MS.
Metals.20 ICP-AES	Determination of various metals by ICP-AES.
Metals.21 CV-AAS	Determination of Mercury by Cold Vapour AAS.
LAB.8	Moisture content determined by heating at 105 deg C for a minimum of 4 hours.
GC.13	Water samples are analysed directly by purge and trap GC-MS.
LAB.4	Toxicity Characteristic Leaching Procedure (TCLP).
EXTRACT.7	Toxicity Characteristic Leaching Procedure (TCLP).
LAB.1	pH - Measured using pH meter and electrode in accordance with APHA 20th ED, 4500-H+.
Metals.20 ICP-AES	Determination of various metals by ICP-AES.
Metals.21 CV-AAS	Determination of Mercury by Cold Vapour AAS.
GC.12 subset	Leachates are extracted with Dichloromethane and analysed by GC-MS.



QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
VOCs in soil						Base II Duplicate II %RPD		
Date extracted	-			29/7/20 10	43953-1	29/7/2010 29/7/2010	LCS-1	29/7/2010
Date analysed	-			31/7/20 10	43953-1	31/7/2010 31/7/2010	LCS-1	31/7/2010
styrene	mg/kg	1	GC.14	<1.0	43953-1	<1.0 <1.0	LCS-1	98%
<i>Surrogate</i> Dibromofluorometha	%		GC.14	95	43953-1	93 90 RPD: 3	LCS-1	97%
Surrogate aaa-Trifluorotoluene	%		GC.14	62	43953-1	70 75 RPD: 7	LCS-1	80%
Surrogate Toluene-da	%		GC.14	105	43953-1	110 109 RPD: 1	LCS-1	98%
Surrogate 4-Bromofluorobenzene	%		GC.14	98	43953-1	94 93 RPD: 1	LCS-1	93%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
vTPH & BTEX in Soil						Base II Duplicate II %RPD		
Date extracted	-			29/7/20 10	43953-1	29/7/2010 29/7/2010	LCS-1	29/7/2010
Date analysed	-			31/7/20 10	43953-1	30/7/2010 30/7/2010	LCS-1	31/7/2010
vTPH C6 - C9	mg/kg	25	GC.16	<25	43953-1	<25 <25	LCS-1	86%
Benzene	mg/kg	0.5	GC.16	<0.5	43953-1	<0.5 <0.5	LCS-1	78%
Toluene	mg/kg	0.5	GC.16	<0.5	43953-1	<0.5 <0.5	LCS-1	88%
Ethylbenzene	mg/kg	1	GC.16	<1.0	43953-1	<1.0 <1.0	LCS-1	88%
m+p-xylene	mg/kg	2	GC.16	<2.0	43953-1	<2.0 <2.0	LCS-1	90%
o-Xylene	mg/kg	1	GC.16	<1.0	43953-1	<1.0 <1.0	LCS-1	92%
Surrogate aaa-Trifluorotoluene	%		GC.16	62	43953-1	70 75 RPD: 7	LCS-1	106%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
sTPH in Soil (C10-C36)						Base II Duplicate II %RPD		
Date extracted	-			29/7/20 10	43953-1	29/7/2010 29/7/2010	LCS-5	29/7/2010
Date analysed	-			29/7/20 10	43953-1	29/7/2010 29/7/2010	LCS-5	29/7/2010
TPH C10 - C14	mg/kg	50	GC.3	<50	43953-1	<50 <50	LCS-5	81%
TPH C15 - C28	mg/kg	100	GC.3	<100	43953-1	320 360 RPD: 12	LCS-5	89%
TPH C29 - C36	mg/kg	100	GC.3	<100	43953-1	380 380 RPD: 0	LCS-5	99%
<i>Surrogate</i> o-Terphenyl	%		GC.3	110	43953-1	# #	LCS-5	96%



Client Reference:

E10100 Macdonaldtown Gasworks

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
PAHs in Soil						Base II Duplicate II %RPD		
Date extracted	-			29/07/2 010	43953-1	29/07/2010 29/07/2010	LCS-5	29/07/2010
Date analysed	-			30/07/2 010	43953-1	30/07/2010 30/07/2010	LCS-5	30/07/2010
Naphthalene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	0.4 0.5 RPD: 22	LCS-5	85%
Acenaphthylene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	0.7 0.9 RPD: 25	[NR]	[NR]
Acenaphthene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	<0.1 0.3	[NR]	[NR]
Fluorene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	0.2 0.4 RPD: 67	LCS-5	102%
Phenanthrene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	2.0 4.9 RPD: 84	LCS-5	92%
Anthracene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	0.6 1.2 RPD: 67	[NR]	[NR]
Fluoranthene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	4.3 7.7 RPD: 57	LCS-5	87%
Pyrene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	5.0 8.1 RPD: 47	LCS-5	108%
Benzo(a)anthracene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	3.3 4.2 RPD: 24	[NR]	[NR]
Chrysene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	2.7 4.2 RPD: 43	LCS-5	108%
Benzo(b+k)fluoranthene	mg/kg	0.2	GC.12 subset	<0.2	43953-1	6.0 6.7 RPD: 11	[NR]	[NR]
Benzo(a)pyrene	mg/kg	0.05	GC.12 subset	<0.05	43953-1	4.4 5.0 RPD: 13	LCS-5	103%
Indeno(1,2,3-c,d)pyrene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	2.0 2.3 RPD: 14	[NR]	[NR]
Dibenzo(a,h)anthracene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	0.5 0.5 RPD: 0	[NR]	[NR]
Benzo(g,h,i)perylene	mg/kg	0.1	GC.12 subset	<0.1	43953-1	1.9 2.2 RPD: 15	[NR]	[NR]
Surrogate p-Terphenyl-d14	%		GC.12 subset	89	43953-1	75 78 RPD: 4	LCS-5	86%

Client Reference: E10100 Mac

E10100 Macdonaldtown Gasworks

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Speciated Phenols in Soil						Base II Duplicate II %RPD		-
Date extracted	-			29/7/20 10	43953-1	29/7/2010 29/7/2010	LCS-1	29/7/2010
Date analysed	-			2/8/201 0	43953-1	2/8/2010 2/8/2010	LCS-1	2/8/2010
Phenol	mg/kg	1	GC.12	<1.0	43953-1	<1.0 <1.0	LCS-1	95%
2-Chlorophenol	mg/kg	1	GC.12	<1.0	43953-1	<1.0 <1.0	LCS-1	102%
2-Methylphenol	mg/kg	1	GC.12	<1.0	43953-1	<1.0 <1.0	[NR]	[NR]
3/4-Methylphenol	mg/kg	2	GC.12	<2.0	43953-1	<2.0 <2.0	[NR]	[NR]
2-Nitrophenol	mg/kg	1	GC.12	<1.0	43953-1	<1.0 <1.0	[NR]	[NR]
2,4-Dimethylphenol	mg/kg	1	GC.12	<1.0	43953-1	<1.0 <1.0	[NR]	[NR]
2,4-Dichlorophenol	mg/kg	1	GC.12	<1.0	43953-1	<1.0 <1.0	[NR]	[NR]
2,6-dichlorophenol	mg/kg	1	GC.12	<1.0	43953-1	<1.0 <1.0	[NR]	[NR]
2,4,5-trichlorophenol	mg/kg	1	GC.12	<1.0	43953-1	<1.0 <1.0	[NR]	[NR]
2,4,6-trichlorophenol	mg/kg	1	GC.12	<1.0	43953-1	<1.0 <1.0	[NR]	[NR]
2,4-dinitrophenol	mg/kg	10	GC.12	<10	43953-1	<10 <10	[NR]	[NR]
4-nitrophenol	mg/kg	10	GC.12	<10	43953-1	<10 <10	LCS-1	74%
2,3,4,6-tetrachlorophenol	mg/kg	1	GC.12	<1.0	43953-1	<1.0 <1.0	[NR]	[NR]
2-methyl-4,6-dinitrophen ol	mg/kg	10	GC.12	<10	43953-1	<10 <10	[NR]	[NR]
pentachlorophenol	mg/kg	10	GC.12	<10	43953-1	<10 <10	[NR]	[NR]
Surrogate 2-fluorophenol	%		GC.12	123	43953-1	105 113 RPD: 7	LCS-1	118%
Surrogate Phenol-d6	%		GC.12	85	43953-1	68 83 RPD: 20	LCS-1	99%
Surrogate 2,4,6-Tribromophenol	%		GC.12	59	43953-1	93 96 RPD: 3	LCS-1	65%
Surrogate p-Terphenyl-d ₁₄	%		GC.12	102	43953-1	114 106 RPD: 7	LCS-1	92%



QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Acid Extractable metals in soil						Base II Duplicate II %RPD		-
Date digested	-	T		29/07/2 010	43953-1	29/07/2010 29/07/2010	LCS-2	29/07/2010
Date analysed	-			29/07/2 010	43953-1	29/07/2010 29/07/2010	LCS-2	29/07/2010
Arsenic	mg/kg	4	Metals.20 ICP-AES	<4	43953-1	36 30 RPD: 18	LCS-2	100%
Cadmium	mg/kg	0.5	Metals.20 ICP-AES	<0.5	43953-1	1.1 1.1 RPD: 0	LCS-2	96%
Chromium	mg/kg	1	Metals.20 ICP-AES	<1	43953-1	19 14 RPD: 30	LCS-2	101%
Lead	mg/kg	1	Metals.20 ICP-AES	<1	43953-1	230 180 RPD: 24	LCS-2	100%
Mercury	mg/kg	0.1	Metals.21 CV-AAS	<0.1	43953-1	0.4 0.3 RPD: 29	LCS-2	97%
Nickel	mg/kg	1	Metals.20 ICP-AES	<1	43953-1	21 17 RPD: 21	LCS-2	104%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank
Moisture				
Date prepared	-			29/7/20 10
Date analysed	-			29/7/20 10
Moisture	%	0.1	LAB.8	<0.10

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
VOCs in Zero Headspace						Base II Duplicate II %RPD		
Date extracted	-			29/7/20 10	[NT]	[NT]	LCS-W1	29/7/2010
Date analysed	-			31/7/20 10	[NT]	[NT]	LCS-W1	31/7/2010
Styrene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	98%
Surrogate Dibromofluoromethane	%		GC.13	95	[NT]	[NT]	LCS-W1	97%
Surrogate toluene-d8	%		GC.13	105	[NT]	[NT]	LCS-W1	98%
Surrogate 4-BFB	%		GC.13	98	[NT]	[NT]	LCS-W1	93%



QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike %
								Recovery
BTEX in Zero Headspace			_			Base II Duplicate II %RPD		
Date extracted	-			29/7/20 10	[NT]	[NT]	LCS-W1	29/7/2010
Date analysed	-			31/7/20 10	[NT]	[NT]	LCS-W1	31/7/2010
Benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	115%
Toluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	106%
Ethylbenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	97%
m+p-xylene	µg/L	2	GC.13	<2.0	[NT]	[NT]	LCS-W1	96%
o-xylene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	98%
<i>Surrogate</i> Dibromofluoromethane	%		GC.13	95	[NT]	[NT]	LCS-W1	102%
Surrogate toluene-d8	%		GC.13	105	[NT]	[NT]	LCS-W1	101%
Surrogate 4-BFB	%		GC.13	98	[NT]	[NT]	LCS-W1	109%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Metals in TCLP USEPA1311						Base II Duplicate II %RPD		
Date extracted	-			29/07/2 010	[NT]	[NT]	LCS-W1	29/07/2010
Date analysed	-			29/07/2 010	[NT]	[NT]	LCS-W1	29/07/2010
Arsenic in TCLP	mg/L	0.05	Metals.20 ICP-AES	<0.05	[NT]	[NT]	LCS-W1	107%
Cadmium in TCLP	mg/L	0.01	Metals.20 ICP-AES	<0.01	[NT]	[NT]	LCS-W1	97%
Chromium in TCLP	mg/L	0.01	Metals.20 ICP-AES	<0.01	[NT]	[NT]	LCS-W1	101%
Lead in TCLP	mg/L	0.03	Metals.20 ICP-AES	<0.03	[NT]	[NT]	LCS-W1	98%
Mercury in TCLP	mg/L	0.0005	Metals.21 CV-AAS	<0.000 5	[NT]	[NT]	LCS-W1	100%
Nickel in TCLP	mg/L	0.02	Metals.20 ICP-AES	<0.02	[NT]	[NT]	LCS-W1	101%



QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
TPH in TCLP extract						Base II Duplicate II %RPD		
Date extracted	-			29/7/20 10	[NT]	[NT]	LCS-W2	29/7/2010
Date analysed	-			29/7/20 10	[NT]	[NT]	LCS-W2	29/7/2010
TPH C10 - C14	µg/L	50	GC.3	<50	[NT]	[NT]	LCS-W2	71%
TPH C15 - C28	µg/L	100	GC.3	<100	[NT]	[NT]	LCS-W2	113%
TPH C29 - C36	µg/L	100	GC.3	<100	[NT]	[NT]	LCS-W2	96%
Surrogate o-Terphenyl	%		GC.3	123	[NT]	[NT]	LCS-W2	116%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
PAHs in TCLP (USEPA 1311)						Base II Duplicate II %RPD		
Date extracted	-			29/07/2 010	[NT]	[NT]	LCS-W1	29/07/2010
Date analysed	-			29/07/2 010	[NT]	[NT]	LCS-W1	29/07/2010
Naphthalene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	91%
Acenaphthylene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Acenaphthene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Fluorene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	108%
Phenanthrene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	103%
Anthracene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Fluoranthene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	99%
Pyrene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	109%
Benzo(a)anthracene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Chrysene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	106%
Benzo(b+k)fluoranthene in TCLP	mg/L	0.002	GC.12 subset	<0.002	[NT]	[NT]	[NR]	[NR]
Benzo(a)pyrene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	LCS-W1	119%
Indeno(1,2,3-c,d)pyrene - TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Dibenzo(a,h)anthracene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
Benzo(g,h,i)perylene in TCLP	mg/L	0.001	GC.12 subset	<0.001	[NT]	[NT]	[NR]	[NR]
<i>Surrogate</i> p-Terphenyl-d ₁₄	%		GC.12	111	[NT]	[NT]	LCS-W1	120%

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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	f Spike % Recovery
Speciated Phenols in water						Base II Duplicate II %RPD		
Date extracted	-			29/07/2 010	[NT]	[NT]	LCS-W	/1 29/07/2010
Date analysed	-			30/07/2 010	[NT]	[NT]	LCS-W ²	/1 30/07/2010
Phenol	µg/L	10	GC.12	<10	[NT]	[NT]	LCS-W	/1 38%
2-Chlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	LCS-W	/1 78%
2-Methylphenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
3/4-Methylphenol	µg/L	20	GC.12	<20	[NT]	[NT]	[NR]	[NR]
2-Nitrophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4-Dimethylphenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4-Dichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,6-Dichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4,5-Trichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4,6-Trichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4-Dinitrophenol	µg/L	100	GC.12	<100	[NT]	[NT]	[NR]	[NR]
4-Nitrophenol	µg/L	100	GC.12	<100	[NT]	[NT]	LCS-W	/1 36%
2,3,4,6-Tetrachlorophen ol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2-methyl-4,6-dinitrophen ol	µg/L	100	GC.12	<100	[NT]	[NT]	[NR]	[NR]
Pentachlorophenol	µg/L	100	GC.12	<100	[NT]	[NT]	[NR]	[NR]
Surrogate 2-fluorophenol	%		GC.12	63	[NT]	[NT]	LCS-W	/1 70%
Surrogate Phenol-d6	%		GC.12	47	[NT]	[NT]	LCS-W	/1 57%
Surrogate 2,4,6-Tribromophenol	%		GC.12	90	[NT]	[NT]	LCS-W ⁷	/1 105%
Surrogate p-Terphenyl-d14	%		GC.12	115	[NT]	[NT]	LCS-W	/1 121%
QUALITY CONTROL vTPH & BTEX in Soil	UNITS	S	Dup. Sm#		Duplicate Duplicate + %RPD	Spike Sm#	Spike % Recov	rery
Date extracted	-	—	[NT]	+	[NT]	43953-2	29/7/2010	
Date analysed	_		[NT]		[NT]	43953-2	31/7/2010	
vTPH C6 - C9	mg/kg		[NT]		[NT]	43953-2	93%	
Benzene							93% 95%	
	mg/kg	-	[NT]		[NT]	43953-2		
Toluene	mg/kg	-	[NT]		[NT]	43953-2	94%	
Ethylbenzene	mg/kg	-	[NT]		[NT]	43953-2	90%	
m+p-xylene	mg/kg	-	[NT]		[NT]	43953-2	92%	
o-Xylene	mg/kថ្	-	[NT]		[NT]	43953-2	95%	
<i>Surrogate</i> aaa-Trifluorotoluene	%		[NT]		[NT]	43953-2	111%	



		Client Reference	e: E10100 Macdonald	Itown Gasworks	5
QUALITY CONTROL sTPH in Soil (C10-C36)	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery
Date extracted	-	[NT]	[NT]	43953-2	29/7/2010
Date analysed	-	[NT]	[NT]	43953-2	29/7/2010
TPH C10 - C14	mg/kg	[NT]	[NT]	43953-2	86%
TPH C15 - C28	mg/kg	[NT]	[NT]	43953-2	#
TPH C29 - C36	mg/kg	[NT]	[NT]	43953-2	#
Surrogate o-Terphenyl	%	[NT]	[NT]	43953-2	#
QUALITY CONTROL PAHs in Soil	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery
Date extracted	-	[NT]	[NT]	43953-2	29/07/2010
Date analysed	-	[NT]	[NT]	43953-2	30/07/2010
Naphthalene	mg/kg	[NT]	[NT]	43953-2	89%
Acenaphthylene	mg/kg	[NT]	[NT]	[NR]	[NR]
Acenaphthene	mg/kg	[NT]	[NT]	[NR]	[NR]
Fluorene	mg/kg	[NT]	[NT]	43953-2	115%
Phenanthrene	mg/kg	[NT]	[NT]	43953-2	#
Anthracene	mg/kg	[NT]	[NT]	[NR]	[NR]
Fluoranthene	mg/kg	[NT]	[NT]	43953-2	#
Pyrene	mg/kg	[NT]	[NT]	43953-2	#
Benzo(a)anthracene	mg/kg	[NT]	[NT]	[NR]	[NR]
Chrysene	mg/kg	[NT]	[NT]	43953-2	#
Benzo(b+k)fluoranthene	mg/kg	[NT]	[NT]	[NR]	[NR]
Benzo(a)pyrene	mg/kg	[NT]	[NT]	43953-2	#
Indeno(1,2,3-c,d)pyrene	mg/kg	[NT]	[NT]	[NR]	[NR]
Dibenzo(a,h)anthracene	mg/kg	[NT]	[NT]	[NR]	[NR]
Benzo(g,h,i)perylene	mg/kg	[NT]	[NT]	[NR]	[NR]
Surrogate p-Terphenyl-d14	%	[NT]	[NT]	43953-2	89%



		Client Reference	6		
QUALITY CONTROL Speciated Phenols in Soil	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery
			Base + Duplicate + %RPD		
Date extracted	-	[NT]	[NT]	43953-2	29/7/2010
Date analysed	-	[NT]	[NT]	43953-2	2/8/2010
Phenol	mg/kg	[NT]	[NT]	43953-2	83%
2-Chlorophenol	mg/kg	[NT]	[NT]	43953-2	90%
2-Methylphenol	mg/kg	[NT]	[NT]	[NR]	[NR]
3/4-Methylphenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2-Nitrophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2,4-Dimethylphenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2,4-Dichlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2,6-dichlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2,4,5-trichlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2,4,6-trichlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2,4-dinitrophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
4-nitrophenol	mg/kg	[NT]	[NT]	43953-2	80%
2,3,4,6-tetrachlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2-methyl-4,6-dinitrophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
pentachlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
Surrogate 2-fluorophenol	%	[NT]	[NT]	43953-2	105%
Surrogate Phenol-d6	%	[NT]	[NT]	43953-2	84%
Surrogate 2,4,6-Tribromophenol	%	[NT]	[NT]	43953-2	107%
<i>Surrogate</i> p-Terphenyl-d ₁₄	%	[NT]	[NT]	43953-2	108%



		Client Reference	e: E10100 Macdonale	dtown Gasworks	
QUALITY CONTROL	UNITS	Dup. Sm#	Duplicate	Spike Sm#	Spike % Recovery
Acid Extractable metals in soil			Base + Duplicate + %RPD		
Date digested	-	[NT]	[NT]	43953-2	29/07/2010
Date analysed	-	[NT]	[NT]	43953-2	29/07/2010
Arsenic	mg/kg	[NT]	[NT]	43953-2	96%
Cadmium	mg/kg	[NT]	[NT]	43953-2	83%
Chromium	mg/kg	[NT]	[NT]	43953-2	92%
Lead	mg/kg	[NT]	[NT]	43953-2	83%
Mercury	mg/kg	[NT]	[NT]	43953-2	92%
Nickel	mg/kg	[NT]	[NT]	43953-2	83%



Report Comments:

Total Petroleum Hydrocarbons in water:# Percent recovery is not possible to report as the high concentration of analytes in the sample/s have caused interference.

Total Petroleum Hydrocarbons in soil:# Percent recovery is not possible to report as the high concentration of analytes in the sample/s have caused interference.

PAH's in soil:# Percent recovery is not possible to report as the high concentration of analytes in the sample/s have caused interference and The RPD for duplicate results is accepted due to the non homogenous national statement of the sample of the non-homogenous national statement of the non-homogenous national statement of the sample of the non-homogenous national statement of the non-homogenous nat

Asbestos was analysed by Approved Ider Asbestos was authorised by Approved Si		Not applicable for this job Not applicable for this job	
INS: Insufficient sample for this test	PQL: Prac	tical Quantitation Limit	NT: Not tested
NA: Test not required	RPD: Rela	tive Percent Difference	NA: Test not required
<: Less than	>: Greater	than	LCS: Laboratory Control Sample

Quality Control Definitions

Blank: This is the component of the analytical signal which is not derived from the sample but from reagents, glassware etc, can be determined by processing solvents and reagents in exactly the same manner as for samples. **Duplicate**: This is the complete duplicate analysis of a sample from the process batch. If possible, the sample selected should be one where the analyte concentration is easily measurable.

Matrix Spike: A portion of the sample is spiked with a known concentration of target analyte. The purpose of the matrix spike is to monitor the performance of the analytical method used and to determine whether matrix interferences exist. **LCS (Laboratory Control Sample)**: This comprises either a standard reference material or a control matrix (such as a blank sand or water) fortified with analytes representative of the analyte class. It is simply a check sample.

Surrogate Spike: Surrogates are known additions to each sample, blank, matrix spike and LCS in a batch, of compounds which are similar to the analyte of interest, however are not expected to be found in real samples.

Laboratory Acceptance Criteria

Duplicate sample and matrix spike recoveries may not be reported on smaller jobs, however, were analysed at a frequency to meet or exceed NEPM requirements. All samples are tested in batched of 20. The duplicate sample RPD and matrix spike recoveries for the batch were within the laboratory acceptance criteria.

Duplicates: <5xPQL - any RPD is acceptable; >5xPQL - 0-50% RPD is acceptable. Matrix Spikes and LCS: Generally 70-130% for inorganics/metals; 60-140% for organics and 10-140% for SVOC and speciated phenols is acceptable.





Envirolab Services Pty Ltd ABN 37 112 535 645 12 Ashley St Chatswood NSW 2067 ph 02 9910 6200 fax 02 9910 6201 enquiries@envirolabservices.com.au www.envirolabservices.com.au

CERTIFICATE OF ANALYSIS 44211

Client: Enviropacific Services (Chatswood) Pty Ltd 1/28 Barcoo St Chatswood NSW 2067

Attention: Marty Croker

Sample log in details:

Your Reference: No. of samples: Date samples received: Date completed instructions received:

E10100 Macdonaldtown Gasworks 7 Soils 02/08/10 02/08/10

Analysis Details:

Please refer to the following pages for results, methodology summary and quality control data. Samples were analysed as received from the client. Results relate specifically to the samples as received. Results are reported on a dry weight basis for solids and on an as received basis for other matrices. *Please refer to the last page of this report for any comments relating to the results.*

Report Details:

 Date results requested by:
 9/08/10

 Date of Preliminary Report:
 Not Issued

 Issue Date:
 9/08/10

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 Accredited for compliance with ISO/IEC 17025.

 Tests not covered by NATA are denoted with *.

Results Approved By:

Sandra Taxlor Senior Organic Chemist

Kluign Morgen

Rhian Morgan Metals Supervisor



Envirolab Reference: 44 Revision No: R

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VOCs in soil						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5%
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	3/8/2010	3/8/2010	3/8/2010	3/8/2010	3/8/2010
Date analysed	-	5/8/2010	5/8/2010	5/8/2010	5/8/2010	5/8/2010
styrene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Surrogate Dibromofluorometha	%	65	69	71	61	69
Surrogate aaa-Trifluorotoluene	%	98	102	89	111	97
Surrogate Toluene-d8	%	90	90	89	89	91
Surrogate 4-Bromofluorobenzene	%	85	85	85	85	89

VOCs in soil			
Our Reference:	UNITS	44211-6	44211-7
Your Reference		Material 3	Control
		Post 20%	
Date Sampled		2/08/2010	2/08/2010
Type of sample		Soil	Soil
Date extracted	-	3/8/2010	3/8/2010
Date analysed	-	5/8/2010	5/8/2010
styrene	mg/kg	<1.0	<1.0
Surrogate Dibromofluorometha	%	70	71
Surrogate aaa-Trifluorotoluene	%	93	85
Surrogate Toluene-da	%	89	88
Surrogate 4-Bromofluorobenzene	%	89	102



vTPH & BTEX in Soil						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5%
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	03/08/2010	03/08/2010	03/08/2010	03/08/2010	03/08/2010
Date analysed	-	05/08/2010	05/08/2010	05/08/2010	05/08/2010	05/08/2010
vTPH C6 - C9	mg/kg	<25	<25	<25	<25	<25
Benzene	mg/kg	<0.5	<0.5	<0.5	<0.5	<0.5
Toluene	mg/kg	<0.5	<0.5	<0.5	<0.5	<0.5
Ethylbenzene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
m+p-xylene	mg/kg	<2.0	<2.0	<2.0	<2.0	<2.0
o-Xylene	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
Surrogate aaa-Trifluorotoluene	%	98	102	89	131	97

vTPH & BTEX in Soil			
Our Reference:	UNITS	44211-6	44211-7
Your Reference		Material 3 Post 20%	Control
Date Sampled		2/08/2010	2/08/2010
Type of sample		Soil	Soil
Date extracted	-	03/08/2010	03/08/2010
Date analysed	-	05/08/2010	05/08/2010
vTPH C6 - C9	mg/kg	<25	<25
Benzene	mg/kg	<0.5	<0.5
Toluene	mg/kg	<0.5	<0.5
Ethylbenzene	mg/kg	<1.0	<1.0
m+p-xylene	mg/kg	<2.0	<2.0
o-Xylene	mg/kg	<1.0	<1.0
Surrogate aaa-Trifluorotoluene	%	93	71

Envirolab Reference: 44211 **Revision No:**



sTPH in Soil (C10-C36)						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5%
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	3/8/2010	3/8/2010	3/8/2010	3/8/2010	3/8/2010
Date analysed	-	3/8/2010	3/8/2010	3/8/2010	3/8/2010	3/8/2010
TPH C10 - C14	mg/kg	<50	<50	<50	<50	<50
TPH C15 - C28	mg/kg	250	200	190	<100	130
TPH C29 - C36	mg/kg	110	120	110	<100	<100
Surrogate o-Terphenyl	%	#	133	131	138	#
sTPH in Soil (C10-C36)				1		

sTPH in Soil (C10-C36)			
Our Reference:	UNITS	44211-6	44211-7
Your Reference		Material 3 Post 20%	Control
Date Sampled		2/08/2010	2/08/2010
Type of sample		Soil	Soil
Date extracted	-	3/8/2010	3/8/2010
Date analysed	-	3/8/2010	3/8/2010
TPH C10 - C14	mg/kg	120	170
TPH C15 - C28	mg/kg	340	330
TPH C29 - C36	mg/kg	<100	<100
Surrogate o-Terphenyl	%	#	#

Envirolab Reference: 44211 **Revision No:**



PAHs in Soil						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5%
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	3/8/2010	3/8/2010	3/8/2010	3/8/2010	3/8/2010
Date analysed	-	5/8/2010	5/8/2010	5/8/2010	5/8/2010	5/8/2010
Naphthalene	mg/kg	0.6	0.5	0.4	2.7	5.0
Acenaphthylene	mg/kg	0.7	0.6	1.4	0.5	0.6
Acenaphthene	mg/kg	<0.1	<0.1	0.2	0.4	0.4
Fluorene	mg/kg	0.2	0.2	0.8	0.9	1.0
Phenanthrene	mg/kg	2.9	2.2	5.6	2.6	2.5
Anthracene	mg/kg	0.8	0.6	1.3	0.7	0.8
Fluoranthene	mg/kg	6.3	4.3	6.6	1.8	1.9
Pyrene	mg/kg	7.1	5.0	6.7	2.3	2.4
Benzo(a)anthracene	mg/kg	4.4	3.3	3.4	0.9	1
Chrysene	mg/kg	4.0	3.0	2.8	0.9	0.9
Benzo(b+k)fluoranthene	mg/kg	6.0	5.0	5.1	1.0	1.1
Benzo(a)pyrene	mg/kg	4.7	3.8	3.9	0.8	0.8
Indeno(1,2,3-c,d)pyrene	mg/kg	2.9	2.2	2.2	0.4	0.5
Dibenzo(a,h)anthracene	mg/kg	0.7	0.6	0.5	0.1	0.1
Benzo(g,h,i)perylene	mg/kg	2.7	2.3	2.2	0.5	0.5
Surrogate p-Terphenyl-d14	%	81	77	76	77	77



Client Reference:

E10100 Macdonaldtown Gasworks

PAHs in Soil			
Our Reference:	UNITS	44211-6	44211-7
Your Reference		Material 3	Control
		Post 20%	
Date Sampled		2/08/2010	2/08/2010
Type of sample		Soil	Soil
Date extracted	-	3/8/2010	3/8/2010
Date analysed	-	5/8/2010	5/8/2010
Naphthalene	mg/kg	16	18
Acenaphthylene	mg/kg	1.2	1.2
Acenaphthene	mg/kg	0.9	0.9
Fluorene	mg/kg	2.8	2.8
Phenanthrene	mg/kg	6.5	6.8
Anthracene	mg/kg	1.8	1.8
Fluoranthene	mg/kg	3.9	4.2
Pyrene	mg/kg	4.9	5.3
Benzo(a)anthracene	mg/kg	1.9	2.0
Chrysene	mg/kg	1.7	1.8
Benzo(b+k)fluoranthene	mg/kg	2.1	2.1
Benzo(a)pyrene	mg/kg	1.7	1.7
Indeno(1,2,3-c,d)pyrene	mg/kg	0.8	0.8
Dibenzo(a,h)anthracene	mg/kg	0.3	0.2
Benzo(g,h,i)perylene	mg/kg	1	1
Surrogate p-Terphenyl-d14	%	77	77

Client Reference:	E10100 Macdonaldtown Gasworks
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Speciated Phenols in Soil						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5%
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	06/08/2010	06/08/2010	06/08/2010	06/08/2010	06/08/2010
Date analysed	-	07/08/2010	07/08/2010	07/08/2010	07/08/2010	07/08/2010
Phenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2-Chlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2-Methylphenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
3/4-Methylphenol	mg/kg	<2.0	<2.0	<2.0	<2.0	<2.0
2-Nitrophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,4-Dimethylphenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,4-Dichlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,6-dichlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,4,5-trichlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,4,6-trichlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2,4-dinitrophenol	mg/kg	<10	<10	<10	<10	<10
4-nitrophenol	mg/kg	<10	<10	<10	<10	<10
2,3,4,6-tetrachlorophenol	mg/kg	<1.0	<1.0	<1.0	<1.0	<1.0
2-methyl-4,6-dinitrophenol	mg/kg	<10	<10	<10	<10	<10
pentachlorophenol	mg/kg	<10	<10	<10	<10	<10
Surrogate 2-fluorophenol	%	#	#	#	82	#
Surrogate Phenol-d6	%	#	#	#	36	#
Surrogate 2,4,6-Tribromophenol	%	#	#	#	18	#
Surrogate p-Terphenyl-d14	%	89	95	97	88	90

E10100 Macdonaldtown Gasworks

Speciated Phenols in Soil			
Our Reference:	UNITS	44211-6	44211-7
Your Reference		Material 3	Control
		Post 20%	
Date Sampled		2/08/2010	2/08/2010
Type of sample		Soil	Soil
Date extracted	-	06/08/2010	06/08/2010
Date analysed	-	07/08/2010	07/08/2010
Phenol	mg/kg	<1.0	<1.0
2-Chlorophenol	mg/kg	<1.0	<1.0
2-Methylphenol	mg/kg	<1.0	<1.0
3/4-Methylphenol	mg/kg	<2.0	<2.0
2-Nitrophenol	mg/kg	<1.0	<1.0
2,4-Dimethylphenol	mg/kg	<1.0	<1.0
2,4-Dichlorophenol	mg/kg	<1.0	<1.0
2,6-dichlorophenol	mg/kg	<1.0	<1.0
2,4,5-trichlorophenol	mg/kg	<1.0	<1.0
2,4,6-trichlorophenol	mg/kg	<1.0	<1.0
2,4-dinitrophenol	mg/kg	<10	<10
4-nitrophenol	mg/kg	<10	<10
2,3,4,6-tetrachlorophenol	mg/kg	<1.0	<1.0
2-methyl-4,6-dinitrophenol	mg/kg	<10	<10
pentachlorophenol	mg/kg	<10	<10
Surrogate 2-fluorophenol	%	#	82
Surrogate Phenol-de	%	#	22
Surrogate 2,4,6-Tribromophenol	%	#	86
Surrogate p-Terphenyl-d14	%	91	109

Acid Extractable metals in soil						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5%
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date digested	-	03/08/2010	03/08/2010	03/08/2010	03/08/2010	03/08/2010
Date analysed	-	04/08/2010	04/08/2010	04/08/2010	04/08/2010	04/08/2010
Arsenic	mg/kg	26	35	29	4	5
Cadmium	mg/kg	1	1.2	1.0	<0.5	<0.5
Chromium	mg/kg	29	21	24	20	21
Lead	mg/kg	170	210	300	16	17
Mercury	mg/kg	0.4	0.4	0.3	<0.1	<0.1
Nickel	mg/kg	20	19	20	2	4

Acid Extractable metals in soil			
Our Reference:	UNITS	44211-6	44211-7
Your Reference		Material 3	Control
		Post 20%	
Date Sampled		2/08/2010	2/08/2010
Type of sample		Soil	Soil
Date digested	-	03/08/2010	03/08/2010
Date analysed	-	04/08/2010	04/08/2010
Arsenic	mg/kg	7	4
Cadmium	mg/kg	<0.5	<0.5
Chromium	mg/kg	28	18
Lead	mg/kg	21	16
Mercury	mg/kg	<0.1	<0.1
Nickel	mg/kg	4	1



Moisture						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5%
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date prepared	-	3/8/2010	3/8/2010	3/8/2010	3/8/2010	3/8/2010
Date analysed	-	3/8/2010	3/8/2010	3/8/2010	3/8/2010	3/8/2010
Moisture	%	12	14	10	23	20
		1		1		
Moisture						
Our Reference:	UNITS	44211-6	44211-7			
Your Reference		Material 3 Post 20%	Control			
Date Sampled		2/08/2010	2/08/2010			
Type of sample		Soil	Soil			
Date prepared	-	3/8/2010	3/8/2010			
Date analysed	-	3/8/2010	3/8/2010			
Moisture	%	19	24			

Envirolab Reference: 44211 **Revision No:**

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VOCs in Zero Headspace ASLP						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5%
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	5/8/2010	5/8/2010	5/8/2010	5/8/2010	5/8/2010
Date analysed	-	5/8/2010	5/8/2010	5/8/2010	5/8/2010	5/8/2010
Styrene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Surrogate Dibromofluoromethane	%	88	81	76	99	87
Surrogate toluene-d8	%	100	99	99	105	109
	%	92	94	90	113	100

VOCs in Zero Headspace ASLP			
Our Reference:	UNITS	44211-6	44211-7
Your Reference		Material 3 Post 20%	Control
Date Sampled		2/08/2010	2/08/2010
Type of sample		Soil	Soil
Date extracted	-	5/8/2010	5/8/2010
Date analysed	-	5/8/2010	5/8/2010
Styrene	μg/L	<1.0	<1.0
Surrogate Dibromofluoromethane	%	89	105
Surrogate toluene-d8	%	111	113
Surrogate 4-BFB	%	106	105

Envirolab Reference: 44211 **Revision No:**



BTEX in Zero Headspace						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5%
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	5/8/2010	5/8/2010	5/8/2010	5/8/2010	5/8/2010
Date analysed	-	5/8/2010	5/8/2010	5/8/2010	5/8/2010	5/8/2010
Benzene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Toluene	µg/L	<1.0	<1.0	<1.0	<1.0	<1.0
Ethylbenzene	µg/L	<1.0	<1.0	<1.0	4.4	<1.0
m+p-xylene	µg/L	<2.0	<2.0	<2.0	<2.0	<2.0
o-xylene	µg/L	<1.0	<1.0	<1.0	2.3	<1.0
Surrogate Dibromofluoromethane	%	88	81	76	99	87
Surrogate toluene-d8	%	100	99	99	105	109
Surrogate 4-BFB	%	92	94	90	113	100

BTEX in Zero Headspace			
Our Reference:	UNITS	44211-6	44211-7
Your Reference		Material 3	Control
		Post 20%	
Date Sampled		2/08/2010	2/08/2010
Type of sample		Soil	Soil
Date extracted	-	5/8/2010	5/8/2010
Date analysed	-	5/8/2010	5/8/2010
Benzene	µg/L	<1.0	<1.0
Toluene	µg/L	<1.0	<1.0
Ethylbenzene	µg/L	2.6	1.5
m+p-xylene	µg/L	<2.0	<2.0
o-xylene	µg/L	1.9	1.5
Surrogate Dibromofluoromethane	%	89	105
Surrogate toluene-d8	%	111	113
Surrogate 4-BFB	%	106	105

Metals-ASLP Neutral (ICP-MS)						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	04/08/2010	04/08/2010	04/08/2010	04/08/2010	04/08/201
Date analysed	-	05/08/2010	05/08/2010	05/08/2010	05/08/2010	05/08/201
pH of final Leachate	pH units	12.3	12.4	12.7	11.5	12.5
Arsenic in ASLP	µg/L	2	<1	<1	2	2
Cadmium in ASLP	µg/L	<0.1	<0.1	<0.1	<0.1	<0.1
Chromium in ASLP	µg/L	42	36	28	13	36
Lead in ASLP	µg/L	1	3	6	<1	<1
Mercury in ASLP	µg/L	<0.50	<0.50	<0.50	<0.50	<0.50
Nickel in ASLP	µg/L	4	5	4	<1	1

Metals-ASLP Neutral (ICP-MS)			
Our Reference:	UNITS	44211-6	44211-7
Your Reference		Material 3	Control
		Post 20%	
Date Sampled		2/08/2010	2/08/2010
Type of sample		Soil	Soil
Date extracted	-	04/08/2010	04/08/2010
Date analysed	-	05/08/2010	05/08/2010
pH of final Leachate	pH units	12.4	9.20
Arsenic in ASLP	µg/L	2	<1
Cadmium in ASLP	µg/L	<0.1	<0.1
Chromium in ASLP	µg/L	43	<1
Lead in ASLP	µg/L	<1	<1
Mercury in ASLP	µg/L	<0.50	<0.50
Nickel in ASLP	µg/L	<1	<1

Envirolab Reference: 44211 **Revision No:**



sTPH in water leach						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5%
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	4/8/2010	4/8/2010	4/8/2010	4/8/2010	4/8/2010
Date analysed	-	4/8/2010	4/8/2010	4/8/2010	4/8/2010	4/8/2010
TPH C10 - C14	µg/L	160	100	110	940	880
TPH C15 - C28	µg/L	240	180	180	480	420
TPH C29 - C36	µg/L	<100	<100	<100	<100	<100
Surrogate o-Terphenyl	%	128	111	114	117	131

sTPH in water leach			
Our Reference:	UNITS	44211-6	44211-7
Your Reference		Material 3	Control
		Post 20%	
Date Sampled		2/08/2010	2/08/2010
Type of sample		Soil	Soil
Date extracted	-	4/8/2010	4/8/2010
Date analysed	-	4/8/2010	4/8/2010
TPH C10 - C14	µg/L	1,400	430
TPH C15 - C28	µg/L	550	250
TPH C29 - C36	µg/L	<100	<100
Surrogate o-Terphenyl	%	119	104



Client Reference:	E10100 Macdonaldtown Gasworks
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PAHs in water leach						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5%
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	4/8/2010	4/8/2010	4/8/2010	4/8/2010	4/8/2010
Date analysed	-	5/8/2010	5/8/2010	5/8/2010	5/8/2010	5/8/2010
Naphthalene in ASLP	mg/L	0.001	<0.001	<0.001	0.20	0.27
Acenaphthylene in ASLP	mg/L	<0.001	<0.001	<0.001	0.002	0.001
Acenaphthene in ASLP	mg/L	<0.001	<0.001	<0.001	0.009	0.008
Fluorene in ASLP	mg/L	<0.001	<0.001	<0.001	0.016	0.017
Phenanthrene in ASLP	mg/L	<0.001	<0.001	<0.001	0.021	0.022
Anthracene in ASLP	mg/L	<0.001	<0.001	<0.001	0.004	0.005
Fluoranthene in ASLP	mg/L	<0.001	<0.001	<0.001	0.003	0.003
Pyrene in ASLP	mg/L	<0.001	<0.001	<0.001	0.003	0.003
Benzo(a)anthracene in ASLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Chrysene in ASLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Benzo(b+k)fluoranthene in ASLP	mg/L	<0.002	<0.002	<0.002	<0.002	<0.002
Benzo(a)pyrene in ASLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Indeno(1,2,3-c,d)pyrene - ASLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Dibenzo(a,h)anthracene in ASLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Benzo(g,h,i)perylene in ASLP	mg/L	<0.001	<0.001	<0.001	<0.001	<0.001
Surrogate p-Terphenyl-d14	%	105	88	95	90	99



Client Reference:

E10100 Macdonaldtown Gasworks

PAHs in water leach			
Our Reference:	UNITS	44211-6	44211-7
Your Reference		Material 3	Control
		Post 20%	
Date Sampled		2/08/2010	2/08/2010
Type of sample		Soil	Soil
Date extracted	-	4/8/2010	4/8/2010
Date analysed	-	5/8/2010	5/8/2010
Naphthalene in ASLP	mg/L	0.42	0.059
Acenaphthylene in ASLP	mg/L	0.002	0.001
Acenaphthene in ASLP	mg/L	0.009	0.004
Fluorene in ASLP	mg/L	0.019	0.007
Phenanthrene in ASLP	mg/L	0.022	0.008
Anthracene in ASLP	mg/L	0.004	0.002
Fluoranthene in ASLP	mg/L	0.003	0.001
Pyrene in ASLP	mg/L	0.003	0.001
Benzo(a)anthracene in ASLP	mg/L	<0.001	<0.001
Chrysene in ASLP	mg/L	<0.001	<0.001
Benzo(b+k)fluoranthene in ASLP	mg/L	<0.002	<0.002
Benzo(a)pyrene in ASLP	mg/L	<0.001	<0.001
Indeno(1,2,3-c,d)pyrene - ASLP	mg/L	<0.001	<0.001
Dibenzo(a,h)anthracene in ASLP	mg/L	<0.001	<0.001
Benzo(g,h,i)perylene in ASLP	mg/L	<0.001	<0.001
Surrogate p-Terphenyl-d14	%	98	90

Client Reference:	E10100 Macdonaldtown Gasworks
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Speciated Phenols in water						
Our Reference:	UNITS	44211-1	44211-2	44211-3	44211-4	44211-5
Your Reference		Material 1	Material 1	Material 1	Material 3	Material 3
		Post 5%	Post 12.5%	Post 20%	Post 5%	Post 12.5%
Date Sampled		2/08/2010	2/08/2010	2/08/2010	2/08/2010	2/08/2010
Type of sample		Soil	Soil	Soil	Soil	Soil
Date extracted	-	4/8/2010	4/8/2010	4/8/2010	4/8/2010	4/8/2010
Date analysed	-	5/8/2010	5/8/2010	5/8/2010	5/8/2010	5/8/2010
Phenol	μg/L	<10	<10	<10	<10	<10
2-Chlorophenol	μg/L	<10	<10	<10	<10	<10
2-Methylphenol	μg/L	<10	<10	<10	<10	<10
3/4-Methylphenol	μg/L	<20	<20	<20	<20	<20
2-Nitrophenol	μg/L	<10	<10	<10	<10	<10
2,4-Dimethylphenol	μg/L	<10	<10	<10	<10	<10
2,4-Dichlorophenol	μg/L	<10	<10	<10	<10	<10
2,6-Dichlorophenol	μg/L	<10	<10	<10	<10	<10
2,4,5-Trichlorophenol	μg/L	<10	<10	<10	<10	<10
2,4,6-Trichlorophenol	µg/L	<10	<10	<10	<10	<10
2,4-Dinitrophenol	µg/L	<100	<100	<100	<100	<100
4-Nitrophenol	µg/L	<100	<100	<100	<100	<100
2,3,4,6-Tetrachlorophenol	µg/L	<10	<10	<10	<10	<10
2-methyl-4,6-dinitrophenol	µg/L	<100	<100	<100	<100	<100
Pentachlorophenol	µg/L	<100	<100	<100	<100	<100
Surrogate 2-fluorophenol	%	62	56	65	74	70
Surrogate Phenol-d6	%	42	38	43	52	47
Surrogate 2,4,6-Tribromophenol	%	129	97	104	127	118
Surrogate p-Terphenyl-d14	%	140	109	121	132	140



Client Reference:	erence:
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E10100 Macdonaldtown Gasworks

Speciated Phenols in water			
Our Reference:	UNITS	44211-6	44211-7
Your Reference		Material 3	Control
		Post 20%	
Date Sampled		2/08/2010	2/08/2010
Type of sample		Soil	Soil
Date extracted	-	4/8/2010	4/8/2010
Date analysed	-	5/8/2010	5/8/2010
Phenol	µg/L	<10	<10
2-Chlorophenol	µg/L	<10	<10
2-Methylphenol	µg/L	<10	<10
3/4-Methylphenol	μg/L	<20	<20
2-Nitrophenol	µg/L	<10	<10
2,4-Dimethylphenol	µg/L	<10	<10
2,4-Dichlorophenol	μg/L	<10	<10
2,6-Dichlorophenol	µg/L	<10	<10
2,4,5-Trichlorophenol	μg/L	<10	<10
2,4,6-Trichlorophenol	μg/L	<10	<10
2,4-Dinitrophenol	µg/L	<100	<100
4-Nitrophenol	µg/L	<100	<100
2,3,4,6-Tetrachlorophenol	µg/L	<10	<10
2-methyl-4,6-dinitrophenol	µg/L	<100	<100
Pentachlorophenol	µg/L	<100	<100
Surrogate 2-fluorophenol	%	83	61
Surrogate Phenol-d6	%	56	34
Surrogate 2,4,6-Tribromophenol	%	110	102
Surrogate p-Terphenyl-d14	%	122	114

Method ID	Methodology Summary
GC.14	Soil samples are extracted with methanol and spiked into water prior to analysing by purge and trap GC-MS.
GC.16	Soil samples are extracted with methanol and spiked into water prior to analysing by purge and trap GC-MS. Water samples are analysed directly by purge and trap GC-MS.
GC.3	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-FID.
GC.12 subset	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-MS.
GC.12	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-MS.
Metals.20 ICP-AES	Determination of various metals by ICP-AES.
Metals.21 CV-AAS	Determination of Mercury by Cold Vapour AAS.
LAB.8	Moisture content determined by heating at 105 deg C for a minimum of 4 hours.
GC.13	Water samples are analysed directly by purge and trap GC-MS.
LAB.1	pH - Measured using pH meter and electrode in accordance with APHA 20th ED, 4500-H+.
Metals.22 ICP-MS	Determination of various metals by ICP-MS following leaching using neutralised deionised water by AS 4439.3 - 1997.
Metals.21 ASLP	Determination of Mercury by Cold Vapour AAS following neutral water leaching by AS 4439.3 - 1997.
GC.12 ASLP	ASLP Leachates are extracted with Dichloromethane and analysed by GC-MS.



QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
VOCs in soil						Base II Duplicate II %RPD		
Date extracted	-			3/8/201 0	44211-1	3/8/2010 3/8/2010	LCS-5	3/8/2010
Date analysed	-			5/8/201 0	44211-1	5/8/2010 5/8/2010	LCS-5	5/8/2010
styrene	mg/kg	1	GC.14	<1.0	44211-1	<1.0 <1.0	LCS-5	110%
<i>Surrogate</i> Dibromofluorometha	%		GC.14	69	44211-1	65 65 RPD: 0	LCS-5	71%
Surrogate aaa-Trifluorotoluene	%		GC.14	106	44211-1	98 108 RPD: 10	LCS-5	113%
Surrogate Toluene-da	%		GC.14	89	44211-1	90 89 RPD: 1	LCS-5	89%
Surrogate 4-Bromofluorobenzene	%		GC.14	84	44211-1	85 84 RPD: 1	LCS-5	83%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
vTPH & BTEX in Soil						Base II Duplicate II %RPD		
Date extracted	-			03/08/2 010	44211-1	03/08/2010 03/08/2010	LCS-1	03/08/2010
Date analysed	-			05/08/2 010	44211-1	05/08/2010 05/08/2010	LCS-1	05/08/2010
vTPH C6 - C9	mg/kg	25	GC.16	<25	44211-1	<25 <25	LCS-1	115%
Benzene	mg/kg	0.5	GC.16	<0.5	44211-1	<0.5 <0.5	LCS-1	72%
Toluene	mg/kg	0.5	GC.16	<0.5	44211-1	<0.5 <0.5	LCS-1	105%
Ethylbenzene	mg/kg	1	GC.16	<1.0	44211-1	<1.0 <1.0	LCS-1	126%
m+p-xylene	mg/kg	2	GC.16	<2.0	44211-1	<2.0 <2.0	LCS-1	136%
o-Xylene	mg/kg	1	GC.16	<1.0	44211-1	<1.0 <1.0	LCS-1	137%
<i>Surrogate</i> aaa-Trifluorotoluene	%		GC.16	106	44211-1	98 137 RPD: 33	LCS-1	105%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
sTPH in Soil (C10-C36)						Base II Duplicate II %RPD		
Date extracted	-			3/8/201 0	44211-1	3/8/2010 3/8/2010	LCS-5	3/8/2010
Date analysed	-			3/8/201 0	44211-1	3/8/2010 3/8/2010	LCS-5	3/8/2010
TPH C10 - C14	mg/kg	50	GC.3	<50	44211-1	<50 <50	LCS-5	76%
TPH C15 - C28	mg/kg	100	GC.3	<100	44211-1	250 200 RPD: 22	LCS-5	81%
TPH C29 - C36	mg/kg	100	GC.3	<100	44211-1	110 130 RPD: 17	LCS-5	80%
Surrogate o-Terphenyl	%		GC.3	127	44211-1	# 134	LCS-5	113%



Client Reference:

E10100 Macdonaldtown Gasworks

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
PAHs in Soil						Base II Duplicate II %RPD		
Date extracted	-			3/8/201 0	44211-1	3/8/2010 3/8/2010	LCS-5	3/8/2010
Date analysed	-			5/8/201 0	44211-1	5/8/2010 5/8/2010	LCS-5	5/8/2010
Naphthalene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	0.6 0.7 RPD: 15	LCS-5	91%
Acenaphthylene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	0.7 0.7 RPD: 0	[NR]	[NR]
Acenaphthene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	<0.1 <0.1	[NR]	[NR]
Fluorene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	0.2 0.3 RPD: 40	LCS-5	95%
Phenanthrene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	2.9 2.6 RPD: 11	LCS-5	103%
Anthracene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	0.8 0.7 RPD: 13	[NR]	[NR]
Fluoranthene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	6.3 4.8 RPD: 27	LCS-5	94%
Pyrene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	7.1 5.4 RPD: 27	LCS-5	101%
Benzo(a)anthracene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	4.4 3.3 RPD: 29	[NR]	[NR]
Chrysene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	4.0 3.1 RPD: 25	LCS-5	112%
Benzo(b+k)fluoranthene	mg/kg	0.2	GC.12 subset	<0.2	44211-1	6.0 5.1 RPD: 16	[NR]	[NR]
Benzo(a)pyrene	mg/kg	0.05	GC.12 subset	<0.05	44211-1	4.7 4.0 RPD: 16	LCS-5	94%
Indeno(1,2,3-c,d)pyrene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	2.9 2.4 RPD: 19	[NR]	[NR]
Dibenzo(a,h)anthracene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	0.7 0.6 RPD: 15	[NR]	[NR]
Benzo(g,h,i)perylene	mg/kg	0.1	GC.12 subset	<0.1	44211-1	2.7 2.4 RPD: 12	[NR]	[NR]
Surrogate p-Terphenyl-d14	%		GC.12 subset	76	44211-1	81 77 RPD: 5	LCS-5	102%

Client Reference: E10100 Macdor

E10100 Macdonaldtown Gasworks

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Speciated Phenols in Soil						Base II Duplicate II %RPD		
Date extracted	-			06/08/2 010	44211-1	06/08/2010 06/08/2010	LCS-5	06/08/2010
Date analysed	-			07/08/2 010	44211-1	07/08/2010 07/08/2010	LCS-5	07/08/2010
Phenol	mg/kg	1	GC.12	<1.0	44211-1	<1.0 <1.0	LCS-5	88%
2-Chlorophenol	mg/kg	1	GC.12	<1.0	44211-1	<1.0 <1.0	LCS-5	95%
2-Methylphenol	mg/kg	1	GC.12	<1.0	44211-1	<1.0 <1.0	[NR]	[NR]
3/4-Methylphenol	mg/kg	2	GC.12	<2.0	44211-1	<2.0 <2.0	[NR]	[NR]
2-Nitrophenol	mg/kg	1	GC.12	<1.0	44211-1	<1.0 <1.0	[NR]	[NR]
2,4-Dimethylphenol	mg/kg	1	GC.12	<1.0	44211-1	<1.0 <1.0	[NR]	[NR]
2,4-Dichlorophenol	mg/kg	1	GC.12	<1.0	44211-1	<1.0 <1.0	[NR]	[NR]
2,6-dichlorophenol	mg/kg	1	GC.12	<1.0	44211-1	<1.0 <1.0	[NR]	[NR]
2,4,5-trichlorophenol	mg/kg	1	GC.12	<1.0	44211-1	<1.0 <1.0	[NR]	[NR]
2,4,6-trichlorophenol	mg/kg	1	GC.12	<1.0	44211-1	<1.0 <1.0	[NR]	[NR]
2,4-dinitrophenol	mg/kg	10	GC.12	<10	44211-1	<10 <10	[NR]	[NR]
4-nitrophenol	mg/kg	10	GC.12	<10	44211-1	<10 <10	LCS-5	26%
2,3,4,6-tetrachlorophenol	mg/kg	1	GC.12	<1.0	44211-1	<1.0 <1.0	[NR]	[NR]
2-methyl-4,6-dinitrophen ol	mg/kg	10	GC.12	<10	44211-1	<10 <10	[NR]	[NR]
pentachlorophenol	mg/kg	10	GC.12	<10	44211-1	<10 <10	[NR]	[NR]
Surrogate 2-fluorophenol	%		GC.12	12	44211-1	# #	LCS-5	61%
Surrogate Phenol-d6	%		GC.12	27	44211-1	# #	LCS-5	67%
Surrogate 2,4,6-Tribromophenol	%		GC.12	15	44211-1	# #	LCS-5	19%
Surrogate p-Terphenyl-d ₁₄	%		GC.12	76	44211-1	89 89 RPD: 0	LCS-5	75%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Acid Extractable metals in soil						Base II Duplicate II %RPD		
Date digested	-	T		03/08/2 010	44211-1	03/08/2010 03/08/2010	LCS-1	03/08/2010
Date analysed	-			04/08/2 010	44211-1	04/08/2010 04/08/2010	LCS-1	04/08/2010
Arsenic	mg/kg	4	Metals.20 ICP-AES	<4	44211-1	26 39 RPD: 40	LCS-1	104%
Cadmium	mg/kg	0.5	Metals.20 ICP-AES	<0.5	44211-1	1 1.2 RPD: 18	LCS-1	106%
Chromium	mg/kg	1	Metals.20 ICP-AES	<1	44211-1	29 20 RPD: 37	LCS-1	102%
Lead	mg/kg	1	Metals.20 ICP-AES	<1	44211-1	170 230 RPD: 30	LCS-1	102%
Mercury	mg/kg	0.1	Metals.21 CV-AAS	<0.1	44211-1	0.4 0.5 RPD: 22	LCS-1	105%
Nickel	mg/kg	1	Metals.20 ICP-AES	<1	44211-1	20 20 RPD: 0	LCS-1	104%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank
Moisture				
Date prepared	-			3/8/201 0
Date analysed	-			3/8/201 0
Moisture	%	0.1	LAB.8	<0.10

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results Base II Duplicate II %RPD	Spike Sm#	Spike % Recovery
Headspace ASLP								
Date extracted	-			5/8/201 0	[NT]	[NT]	LCS-W1	05/08/2010
Date analysed	-			5/8/201 0	[NT]	[NT]	LCS-W1	05/08/2010
Styrene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	104%
Surrogate Dibromofluoromethane	%		GC.13	98	[NT]	[NT]	LCS-W1	106%
Surrogate toluene-d8	%		GC.13	104	[NT]	[NT]	LCS-W1	103%
Surrogate 4-BFB	%		GC.13	104	[NT]	[NT]	LCS-W1	102%



QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike %
QUALITY CONTROL			METHOD	Dianix	Duplicate on #	Duplicate results		Recovery
BTEX in Zero Headspace						Base II Duplicate II %RPD		
Date extracted	-			5/8/201 0	[NT]	[NT]	LCS-W1	5/8/2010
Date analysed	-			5/8/201 0	[NT]	[NT]	LCS-W1	5/8/2010
Benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Toluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Ethylbenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
m+p-xylene	µg/L	2	GC.13	<2.0	[NT]	[NT]	[NR]	[NR]
o-xylene	µg/L	1	GC.13	<1.0	[NT]	[NT]	[NR]	[NR]
Surrogate Dibromofluoromethane	%		GC.13	98	[NT]	[NT]	LCS-W1	98%
Surrogate toluene-d8	%		GC.13	100	[NT]	[NT]	LCS-W1	100%
Surrogate 4-BFB	%		GC.13	104	[NT]	[NT]	LCS-W1	104%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Metals-ASLP Neutral (ICP-MS)						Base II Duplicate II %RPD		
Date extracted	-	+	1	[NT]	44211-1	04/08/2010 04/08/2010	LCS-W1	05/08/2010
Date analysed	-			05/08/2 010	44211-1	05/08/2010 05/08/2010	[NR]	[NR]
Arsenic in ASLP	µg/L	1	Metals.22 ICP-MS	<1	44211-1	2 2 RPD: 0	LCS-W1	106%
Cadmium in ASLP	µg/L	0.1	Metals.22 ICP-MS	<0.1	44211-1	<0.1 <0.1	LCS-W1	110%
Chromium in ASLP	µg/L	1	Metals.22 ICP-MS	<1	44211-1	42 42 RPD: 0	LCS-W1	102%
Lead in ASLP	µg/L	1	Metals.22 ICP-MS	<1	44211-1	1 1 RPD: 0	LCS-W1	106%
Mercury in ASLP	µg/L	0.5	Metals.21 ASLP	<0.50	44211-1	<0.50 <0.50	LCS-W1	117%
Nickel in ASLP	µg/L	1	Metals.22 ICP-MS	<1	44211-1	4 4 RPD: 0	LCS-W1	98%



QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
sTPH in water leach						Base II Duplicate II %RPD		
Date extracted	-			4/8/201 0	[NT]	[NT]	LCS-W1	4/8/2010
Date analysed	-			4/8/201 0	[NT]	[NT]	LCS-W1	4/8/2010
TPH C10 - C14	µg/L	50	GC.3	<50	[NT]	[NT]	LCS-W1	83%
TPH C15 - C28	µg/L	100	GC.3	<100	[NT]	[NT]	LCS-W1	120%
TPH C29 - C36	µg/L	100	GC.3	<100	[NT]	[NT]	LCS-W1	90%
<i>Surrogate</i> o-Terphenyl	%		GC.3	116	[NT]	[NT]	LCS-W1	112%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
PAHs in water leach						Base II Duplicate II %RPD		,
Date extracted	-			04/08/2 010	[NT]	[NT]	LCS-W1	04/08/2010
Date analysed	-			05/08/2 010	[NT]	[NT]	LCS-W1	05/08/2010
Naphthalene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	92%
Acenaphthylene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Acenaphthene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Fluorene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	110%
Phenanthrene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	109%
Anthracene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Fluoranthene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	104%
Pyrene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	114%
Benzo(a)anthracene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Chrysene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	106%
Benzo(b+k)fluoranthene in ASLP	mg/L	0.002	GC.12 ASLP	<0.002	[NT]	[NT]	[NR]	[NR]
Benzo(a)pyrene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	113%
Indeno(1,2,3-c,d)pyrene - ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Dibenzo(a,h)anthracene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Benzo(g,h,i)perylene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Surrogate p-Terphenyl-d14	%		GC.12	122	[NT]	[NT]	LCS-W1	105%

Envirolab Reference: 44211 Revision No: R 00

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike %
								Recovery
Speciated Phenols in water						Base II Duplicate II %RPD		
Date extracted	-			4/8/201 0	[NT]	[NT]	LCS-W1	4/8/2010
Date analysed	-			5/8/201 0	[NT]	[NT]	LCS-W1	5/8/2010
Phenol	µg/L	10	GC.12	<10	[NT]	[NT]	LCS-W1	27%
2-Chlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	LCS-W1	84%
2-Methylphenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
3/4-Methylphenol	µg/L	20	GC.12	<20	[NT]	[NT]	[NR]	[NR]
2-Nitrophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4-Dimethylphenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4-Dichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,6-Dichlorophenol	μg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4,5-Trichlorophenol	μg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4,6-Trichlorophenol	μg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4-Dinitrophenol	μg/L	100	GC.12	<100	[NT]	[NT]	[NR]	[NR]
4-Nitrophenol	μg/L	100	GC.12	<100	[NT]	[NT]	LCS-W1	28%
2,3,4,6-Tetrachlorophen ol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2-methyl-4,6-dinitrophen ol	µg/L	100	GC.12	<100	[NT]	[NT]	[NR]	[NR]
Pentachlorophenol	µg/L	100	GC.12	<100	[NT]	[NT]	[NR]	[NR]
Surrogate 2-fluorophenol	%		GC.12	60	[NT]	[NT]	LCS-W1	57%
Surrogate Phenol-d6	%		GC.12	31	[NT]	[NT]	LCS-W1	30%
Surrogate 2,4,6-Tribromophenol	%		GC.12	124	[NT]	[NT]	LCS-W1	120%
Surrogate p-Terphenyl-d ₁₄	%		GC.12	127	[NT]	[NT]	LCS-W1	119%
QUALITY CONTROL vTPH & BTEX in Soil	UNITS	S	Dup. Sm#		Duplicate Duplicate + %RPD		Spike % Recovery	
Date extracted	<u> </u>	=	[NT]	+	[NT]	43953-2	03/08/2010	=
Date analysed	_		[NT]		[NT]	43953-2	05/08/2010	
vTPH C6 - C9	mg/kg	a	[NT]		[NT]	43953-2	96%	
Benzene	mg/kg		[NT]		[NT]	43953-2	60%	
		-					87%	
Toluene	mg/ko	-	[NT]		[NT]	43953-2		
Ethylbenzene m+p-xylene	mg/kg mg/kg	-	[NT] [NT]		[NT] [NT]	43953-2 43953-2	103% 114%	
		-					113%	
o-Xylene <i>Surrogate</i> aaa-Trifluorotoluene	mg/kg	9	[NT] [NT]		[NT] [NT]	43953-2 43953-2	91%	



Client Reference: E10100 Macdonaldtown Gasworks								
QUALITY CONTROL sTPH in Soil (C10-C36)	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery			
Date extracted	-	[NT]	[NT]	43953-2	3/8/2010			
Date analysed	-	[NT]	[NT]	43953-2	3/8/2010			
TPH C10 - C14	mg/kg	[NT]	[NT]	43953-2	80%			
TPH C15 - C28	mg/kg	[NT]	[NT]	43953-2	139%			
TPH C29 - C36	mg/kg	[NT]	[NT]	43953-2	99%			
Surrogate o-Terphenyl	%	[NT]	[NT]	43953-2	131%			
QUALITY CONTROL PAHs in Soil	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery			
Date extracted	-	[NT]	[NT]	43953-2	3/8/2010			
Date analysed	-	[NT]	[NT]	43953-2	5/8/2010			
Naphthalene	mg/kg	[NT]	[NT]	43953-2	90%			
Acenaphthylene	mg/kg	[NT]	[NT]	[NR]	[NR]			
Acenaphthene	mg/kg	[NT]	[NT]	[NR]	[NR]			
Fluorene	mg/kg	[NT]	[NT]	43953-2	101%			
Phenanthrene	mg/kg	[NT]	[NT]	43953-2	77%			
Anthracene	mg/kg	[NT]	[NT]	[NR]	[NR]			
Fluoranthene	mg/kg	[NT]	[NT]	43953-2	68%			
Pyrene	mg/kg	[NT]	[NT]	43953-2	70%			
Benzo(a)anthracene	mg/kg	[NT]	[NT]	[NR]	[NR]			
Chrysene	mg/kg	[NT]	[NT]	43953-2	79%			
Benzo(b+k)fluoranthene	mg/kg	[NT]	[NT]	[NR]	[NR]			
Benzo(a)pyrene	mg/kg	[NT]	[NT]	43953-2	91%			
Indeno(1,2,3-c,d)pyrene	mg/kg	[NT]	[NT]	[NR]	[NR]			
Dibenzo(a,h)anthracene	mg/kg	[NT]	[NT]	[NR]	[NR]			
Benzo(g,h,i)perylene	mg/kg	[NT]	[NT]	[NR]	[NR]			
Surrogate p-Terphenyl-d14	%	[NT]	[NT]	43953-2	76%			



		Client Referenc	e: E10100 Macdonald	town Gasworks	5
QUALITY CONTROL Speciated Phenols in Soil	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery
Date extracted	-	[NT]	[NT]	44211-2	06/08/2010
Date analysed	-	[NT]	[NT]	44211-2	07/08/2010
Phenol	mg/kg	[NT]	[NT]	44211-2	32%
2-Chlorophenol	mg/kg	[NT]	[NT]	44211-2	#
2-Methylphenol	mg/kg	[NT]	[NT]	[NR]	[NR]
3/4-Methylphenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2-Nitrophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2,4-Dimethylphenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2,4-Dichlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2,6-dichlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2,4,5-trichlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2,4,6-trichlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2,4-dinitrophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
4-nitrophenol	mg/kg	[NT]	[NT]	44211-2	13%
2,3,4,6-tetrachlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2-methyl-4,6-dinitrophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
pentachlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
<i>Surrogate</i> 2-fluorophenol	%	[NT]	[NT]	44211-2	#
Surrogate Phenol-d6	%	[NT]	[NT]	44211-2	#
Surrogate 2,4,6-Tribromophenol	%	[NT]	[NT]	44211-2	#
Surrogate p-Terphenyl-d14	%	[NT]	[NT]	44211-2	84%

		Client Reference	ce: E10100 Macdonald	ltown Gasworks	
QUALITY CONTROL Acid Extractable metals in soil	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery
Date digested	-	[NT]	[NT]	43953-2	03/08/2010
Date analysed	-	[NT]	[NT]	43953-2	04/08/2010
Arsenic	mg/kg	[NT]	[NT]	43953-2	99%
Cadmium	mg/kg	[NT]	[NT]	43953-2	88%
Chromium	mg/kg	[NT]	[NT]	43953-2	88%
Lead	mg/kg	[NT]	[NT]	43953-2	112%
Mercury	mg/kg	[NT]	[NT]	43953-2	108%
Nickel	mg/kg	[NT]	[NT]	43953-2	89%
QUALITY CONTROL Metals-ASLP Neutral (ICP-MS)	UNITS	Dup. Sm#	Duplicate Base + Duplicate + %RPD	Spike Sm#	Spike % Recovery
Date extracted	-	[NT]	[NT]	44211-2	05/08/2010
Date analysed	-	[NT]	[NT]	[NR]	[NR]
Arsenic in ASLP	µg/L	[NT]	[NT]	44211-2	106%
Cadmium in ASLP	µg/L	[NT]	[NT]	44211-2	108%
Chromium in ASLP	µg/L	[NT]	[NT]	44211-2	102%
Lead in ASLP	µg/L	[NT]	[NT]	44211-2	102%
Mercury in ASLP	µg/L	[NT]	[NT]	44211-2	106%
Nickel in ASLP	µg/L	[NT]	[NT]	44211-2	96%



Report Comments:

Total Petroleum Hydrocarbons in water:# Percent recovery is not possible to report as the high concentration of analytes in the sample/s have caused interference.

Total Petroleum Hydrocarbons in soil:# Percent recovery is not possible to report as the high concentration of analytes in the sample/s have caused interference.

PAH's in soil:# Percent recovery is not possible to report as the high concentration of analytes in the sample/s have caused interference and The RPD for duplicate results is accepted due to the non homogenous nature of the sample/s.

Total Petroleum Hydrocarbons in soil:# Percent recovery is not possible to report as the high concentration of analytes in the sample/s have caused interference.

Phenols in soil:# Percent recovery is not possible to report due to interference from analytes (other than those being tested) in the sample/s.

Asbestos was analysed by Approved Identifier: Not applicable for this job

Asbestos was authorised by Approved Signatory: Not applicable for this job

INS: Insufficient sample for this test NT: Not tested PQL: Practical Quantitation Limit <: Less than >: Greater than RPD: Relative Percent Difference NA: Test not required LCS: Laboratory Control Sample NR: Not requested

Quality Control Definitions

Blank: This is the component of the analytical signal which is not derived from the sample but from reagents, glassware etc, can be determined by processing solvents and reagents in exactly the same manner as for samples. **Duplicate**: This is the complete duplicate analysis of a sample from the process batch. If possible, the sample selected should be one where the analyte concentration is easily measurable.

Matrix Spike: A portion of the sample is spiked with a known concentration of target analyte. The purpose of the matrix spike is to monitor the performance of the analytical method used and to determine whether matrix interferences exist. LCS (Laboratory Control Sample): This comprises either a standard reference material or a control matrix (such as a blank sand or water) fortified with analytes representative of the analyte class. It is simply a check sample.

Surrogate Spike: Surrogates are known additions to each sample, blank, matrix spike and LCS in a batch, of compounds which are similar to the analyte of interest, however are not expected to be found in real samples.

Laboratory Acceptance Criteria:

Duplicate sample and matrix spike recoveries may not be reported on smaller jobs, however, were analysed at a frequency to meet or exceed NEPM requirements. All samples are tested in batches of 20. The duplicate sample RPD and matrix spike recoveries for the sample batch were within laboratory acceptance criteria.

Duplicates: <5xPQL - any RPD is acceptable; >5xPQL - 0-50% RPD is acceptable.

Matrix Spikes and LCS: Generally 70-130% for inorganics/metals; 60-140% for organics and 10-140% for

SVOC and speciated phenols is acceptable. Surrogates: 60-140% is acceptable for general organics and 10-140% for



Envirolab Services Pty Ltd ABN 37 112 535 645 12 Ashley St Chatswood NSW 2067 ph 02 9910 6200 fax 02 9910 6201 enquiries@envirolabservices.com.au www.envirolabservices.com.au

CERTIFICATE OF ANALYSIS 44271

Client: Enviropacific Services (Chatswood) Pty Ltd 1/28 Barcoo St Chatswood NSW 2067

Attention: Marty Croker

Sample log in details:

Your Reference: No. of samples: Date samples received: Date completed instructions received:

E10100 Macdonaldtown Gasworks 3 Soils 04/08/10 04/08/10

Analysis Details:

Please refer to the following pages for results, methodology summary and quality control data. Samples were analysed as received from the client. Results relate specifically to the samples as received. Results are reported on a dry weight basis for solids and on an as received basis for other matrices. *Please refer to the last page of this report for any comments relating to the results.*

Report Details:

 Date results requested by:
 11/08/10

 Date of Preliminary Report:
 Not issued

 Issue Date:
 11/08/10

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 Accredited for compliance with ISO/IEC 17025.

 Tests not covered by NATA are denoted with *.

Results Approved By:

Giovanni Agosti Technical Manager

₹a. Sandra Taxlor Senior Organic Chemist

Kluign Morgen

Rhian Morgan Metals Supervisor

Jacinta Hurst Laboratory Manager

Envirolab Reference: 44271 Revision No: R 00 Nancy Zhang Chemist



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VOCs in soil				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2	Material 2	Material 2
		Post 5%	Post 12.5%	Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date extracted	-	9/8/2010	9/8/2010	9/8/2010
Date analysed	-	10/8/2010	10/8/2010	10/8/2010
styrene	mg/kg	<1.0	<1.0	<1.0
Surrogate Dibromofluorometha	%	96	87	87
Surrogate aaa-Trifluorotoluene	%	70	71	72
Surrogate Toluene-da	%	95	102	104
Surrogate 4-Bromofluorobenzene	%	101	96	96

Envirolab Reference: 44271 **Revision No:**



Client Reference: E10100 Macdonaldtown Gasworks	Client Reference:	E10100 Macdonaldtown Gasworks
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vTPH & BTEX in Soil				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2	Material 2	Material 2
		Post 5%	Post 12.5%	Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date extracted	-	9/8/2010	9/8/2010	9/8/2010
Date analysed	-	10/8/2010	10/8/2010	10/8/2010
vTPH C6 - C9	mg/kg	<25	<25	<25
Benzene	mg/kg	<0.5	<0.5	<0.5
Toluene	mg/kg	<0.5	<0.5	<0.5
Ethylbenzene	mg/kg	2.8	1.4	<1.0
m+p-xylene	mg/kg	2.4	<2.0	<2.0
o-Xylene	mg/kg	2.3	1.2	<1.0
Surrogate aaa-Trifluorotoluene	%	70	71	72



Client Reference: E10100 Mac

E10100 Macdonaldtown Gasworks

sTPH in Soil (C10-C36)				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2	Material 2	Material 2
		Post 5%	Post 12.5%	Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date extracted	-	9/8/2010	9/8/2010	9/8/2010
Date analysed	-	9/8/2010	9/8/2010	9/8/2010
TPH C10 - C14	mg/kg	470	240	210
TPH C15 - C28	mg/kg	340	210	210
TPH C29 - C36	mg/kg	<100	<100	<100
Surrogate o-Terphenyl	%	#	#	#

Client	Reference
0110111	

e: E10100 Macdonaldtown Gasworks

PAHs in Soil				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2	Material 2	Material 2
		Post 5%	Post 12.5%	Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date extracted	-	9/8/2010	9/8/2010	9/8/2010
Date analysed	-	10/8/2010	10/8/2010	10/8/2010
Naphthalene	mg/kg	120	74	60
Acenaphthylene	mg/kg	3.2	2.5	2.4
Acenaphthene	mg/kg	6.9	5.5	5.1
Fluorene	mg/kg	6.2	5.0	4.7
Phenanthrene	mg/kg	12	9.5	9.2
Anthracene	mg/kg	3.2	2.7	2.5
Fluoranthene	mg/kg	5.2	4.2	4.2
Pyrene	mg/kg	6.3	5.2	4.9
Benzo(a)anthracene	mg/kg	2.2	1.8	1.8
Chrysene	mg/kg	2.1	1.7	1.7
Benzo(b+k)fluoranthene	mg/kg	0.7	1.8	1.9
Benzo(a)pyrene	mg/kg	2.0	1.6	1.6
Indeno(1,2,3-c,d)pyrene	mg/kg	0.5	0.5	0.4
Dibenzo(a,h)anthracene	mg/kg	0.2	0.1	0.1
Benzo(g,h,i)perylene	mg/kg	0.6	0.5	0.5
Surrogate p-Terphenyl-d14	%	95	97	99



Client Reference:

E10100 Macdonaldtown Gasworks

Speciated Phenols in Soil				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2	Material 2	Material 2
		Post 5%	Post 12.5%	Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date extracted	-	10/08/2010	10/08/2010	10/08/2010
Date analysed	-	11/08/2010	11/08/2010	11/08/2010
Phenol	mg/kg	<1.0	<1.0	<1.0
2-Chlorophenol	mg/kg	<1.0	<1.0	<1.0
2-Methylphenol	mg/kg	<1.0	<1.0	<1.0
3/4-Methylphenol	mg/kg	<2.0	<2.0	<2.0
2-Nitrophenol	mg/kg	<1.0	<1.0	<1.0
2,4-Dimethylphenol	mg/kg	<1.0	<1.0	<1.0
2,4-Dichlorophenol	mg/kg	<1.0	<1.0	<1.0
2,6-dichlorophenol	mg/kg	<1.0	<1.0	<1.0
2,4,5-trichlorophenol	mg/kg	<1.0	<1.0	<1.0
2,4,6-trichlorophenol	mg/kg	<1.0	<1.0	<1.0
2,4-dinitrophenol	mg/kg	<10	<10	<10
4-nitrophenol	mg/kg	<10	<10	<10
2,3,4,6-tetrachlorophenol	mg/kg	<1.0	<1.0	<1.0
2-methyl-4,6-dinitrophenol	mg/kg	<10	<10	<10
pentachlorophenol	mg/kg	<10	<10	<10
Surrogate 2-fluorophenol	%	64	#	#
Surrogate Phenol-de	%	77	56	43
Surrogate 2,4,6-Tribromophenol	%	#	#	#
Surrogate p-Terphenyl-d14	%	107	102	102

Client Reference:	E10100 Macdonaldtown Gasworks

Acid Extractable metals in soil				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2	Material 2	Material 2
		Post 5%	Post 12.5%	Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date digested	-	09/08/2010	09/08/2010	09/08/2010
Date analysed	-	10/08/2010	10/08/2010	10/08/2010
Arsenic	mg/kg	4	<4	5
Cadmium	mg/kg	<0.5	<0.5	<0.5
Chromium	mg/kg	15	16	21
Lead	mg/kg	24	26	23
Mercury	mg/kg	<0.1	<0.1	<0.1
Nickel	mg/kg	3	4	5

Envirolab Reference: 44271 **Revision No:**



Moisture				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2 Post 5%	Material 2 Post 12.5%	Material 2 Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date prepared	-	9/8/2010	9/8/2010	9/8/2010
Date analysed	-	9/8/2010	9/8/2010	9/8/2010
	%	14	13	12

Envirolab Reference: 44271 **Revision No:**



Client Reference: E

E10100 Macdonaldtown Gasworks

VOCs in Zero Headspace ASLP				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2	Material 2	Material 2
		Post 5%	Post 12.5%	Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date extracted	-	10/8/2010	10/8/2010	10/8/2010
Date analysed	-	10/8/2010	10/8/2010	10/8/2010
Styrene	µg/L	2.1	2.0	1.7
Surrogate Dibromofluoromethane	%	91	89	87
Surrogate toluene-d8	%	95	109	107
Surrogate 4-BFB	%	113	115	106

Client	Reference:	E10100
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E10100 Macdonaldtown Gasworks

BTEX in Zero Headspace				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2	Material 2	Material 2
		Post 5%	Post 12.5%	Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date extracted	-	10/8/2010	10/8/2010	10/8/2010
Date analysed	-	10/8/2010	10/8/2010	10/8/2010
Benzene	µg/L	<1.0	<1.0	<1.0
Toluene	µg/L	<1.0	<1.0	<1.0
Ethylbenzene	µg/L	70	44	30
m+p-xylene	µg/L	62	41	26
o-xylene	μg/L	59	39	26
Surrogate Dibromofluoromethane	%	91	89	87
Surrogate toluene-d8	%	95	109	107
Surrogate 4-BFB	%	113	115	106

Client	Reference:	E10100	Macdon

E10100	Macdonaldtown	Gasworks
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Metals-ASLP Neutral (ICP-MS)				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2	Material 2	Material 2
		Post 5%	Post 12.5%	Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date extracted	-	11/08/2010	11/08/2010	11/08/2010
Date analysed	-	11/08/2010	11/08/2010	11/08/2010
pH of final Leachate	pH units	11.6	12.2	12.5
Arsenic in ASLP	µg/L	8	7	6
Cadmium in ASLP	µg/L	<0.1	<0.1	0.1
Chromium in ASLP	µg/L	7	26	38
Lead in ASLP	µg/L	<1	<1	<1
Mercury in ASLP	µg/L	<0.50	<0.50	<0.50
Nickel in ASLP	μg/L	2	4	5

Envirolab Reference: 44271 **Revision No:**



sTPH in water leach				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2	Material 2	Material 2
		Post 5%	Post 12.5%	Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date extracted	-	5/8/2010	5/8/2010	5/8/2010
Date analysed	-	5/8/2010	5/8/2010	5/8/2010
TPH C10 - C14	µg/L	5,600	4,500	3,700
TPH C15 - C28	µg/L	1,100	1,200	1,200
TPH C29 - C36	µg/L	<100	<100	<100
Surrogate o-Terphenyl	%	#	#	#

Client Reference:

E10100 Macdonaldtown Gasworks

PAHs in water leach				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2	Material 2	Material 2
		Post 5%	Post 12.5%	Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date extracted	-	05/08/2010	05/08/2010	05/08/2010
Date analysed	-	06/08/2010	06/08/2010	06/08/2010
Naphthalene in ASLP	mg/L	2.4	1.7	1.3
Acenaphthylene in ASLP	mg/L	0.024	0.026	0.028
Acenaphthene in ASLP	mg/L	0.054	0.059	0.064
Fluorene in ASLP	mg/L	0.036	0.037	0.042
Phenanthrene in ASLP	mg/L	0.026	0.029	0.034
Anthracene in ASLP	mg/L	0.005	0.006	0.007
Fluoranthene in ASLP	mg/L	0.003	0.003	0.004
Pyrene in ASLP	mg/L	0.002	0.003	0.004
Benzo(a)anthracene in ASLP	mg/L	<0.001	<0.001	<0.001
Chrysene in ASLP	mg/L	<0.001	<0.001	<0.001
Benzo(b+k)fluoranthene in ASLP	mg/L	<0.002	<0.002	<0.002
Benzo(a)pyrene in ASLP	mg/L	<0.001	<0.001	<0.001
Indeno(1,2,3-c,d)pyrene - ASLP	mg/L	<0.001	<0.001	<0.001
Dibenzo(a,h)anthracene in ASLP	mg/L	<0.001	<0.001	<0.001
Benzo(g,h,i)perylene in ASLP	mg/L	<0.001	<0.001	<0.001
Surrogate p-Terphenyl-d14	%	85	87	98

Client Reference:

E10100 Macdonaldtown Gasworks

Speciated Phenols in water				
Our Reference:	UNITS	44271-1	44271-2	44271-3
Your Reference		Material 2 Post 5%	Material 2 Post 12.5%	Material 2 Post 20%
Date Sampled		4/08/2010	4/08/2010	4/08/2010
Type of sample		Soil	Soil	Soil
Date extracted	-	05/08/2010	05/08/2010	05/08/2010
Date analysed	-	05/08/2010	05/08/2010	05/08/2010
Phenol	µg/L	<10	<10	<10
2-Chlorophenol	µg/L	<10	<10	<10
2-Methylphenol	µg/L	<10	<10	<10
3/4-Methylphenol	µg/L	<20	<20	<20
2-Nitrophenol	µg/L	<10	<10	<10
2,4-Dimethylphenol	µg/L	<10	<10	<10
2,4-Dichlorophenol	µg/L	<10	<10	<10
2,6-Dichlorophenol	µg/L	<10	<10	<10
2,4,5-Trichlorophenol	µg/L	<10	<10	<10
2,4,6-Trichlorophenol	µg/L	<10	<10	<10
2,4-Dinitrophenol	µg/L	<100	<100	<100
4-Nitrophenol	µg/L	<100	<100	<100
2,3,4,6-Tetrachlorophenol	µg/L	<10	<10	<10
2-methyl-4,6-dinitrophenol	µg/L	<100	<100	<100
Pentachlorophenol	µg/L	<100	<100	<100
Surrogate 2-fluorophenol	%	65	62	62
Surrogate Phenol-de	%	41	42	42
Surrogate 2,4,6-Tribromophenol	%	117	110	129
Surrogate p-Terphenyl-d14	%	120	98	132

Method ID	Methodology Summary
GC.14	Soil samples are extracted with methanol and spiked into water prior to analysing by purge and trap GC-MS.
GC.16	Soil samples are extracted with methanol and spiked into water prior to analysing by purge and trap GC-MS. Water samples are analysed directly by purge and trap GC-MS.
GC.3	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-FID.
GC.12 subset	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-MS.
GC.12	Soil samples are extracted with Dichloromethane/Acetone and waters with Dichloromethane and analysed by GC-MS.
Metals.20 ICP-AES	Determination of various metals by ICP-AES.
Metals.21 CV-AAS	Determination of Mercury by Cold Vapour AAS.
LAB.8	Moisture content determined by heating at 105 deg C for a minimum of 4 hours.
GC.13	Water samples are analysed directly by purge and trap GC-MS.
LAB.1	pH - Measured using pH meter and electrode in accordance with APHA 20th ED, 4500-H+.
Metals.22 ICP-MS	Determination of various metals by ICP-MS following leaching using neutralised deionised water by AS 4439.3 - 1997.
Metals.21 ASLP	Determination of Mercury by Cold Vapour AAS following neutral water leaching by AS 4439.3 - 1997.
GC.12 ASLP	ASLP Leachates are extracted with Dichloromethane and analysed by GC-MS.



QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
VOCs in soil						Base II Duplicate II %RPD		
Date extracted	-			9/8/201 0	[NT]	[NT]	LCS-5	9/8/2010
Date analysed	-			10/8/20 10	[NT]	[NT]	LCS-5	10/8/2010
styrene	mg/kg	1	GC.14	<1.0	[NT]	[NT]	LCS-5	104%
Surrogate Dibromofluorometha	%		GC.14	96	[NT]	[NT]	LCS-5	86%
Surrogate aaa-Trifluorotoluene	%		GC.14	72	[NT]	[NT]	LCS-5	76%
<i>Surrogate</i> Toluene-dଃ	%		GC.14	93	[NT]	[NT]	LCS-5	94%
Surrogate 4-Bromofluorobenzene	%		GC.14	79	[NT]	[NT]	LCS-5	82%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
vTPH & BTEX in Soil						Base II Duplicate II %RPD		
Date extracted	-			9/8/201 0	[NT]	[NT]	LCS-5	9/8/2010
Date analysed	-			10/8/20 10	[NT]	[NT]	LCS-5	10/8/2010
vTPH C6 - C9	mg/kg	25	GC.16	<25	[NT]	[NT]	LCS-5	87%
Benzene	mg/kg	0.5	GC.16	<0.5	[NT]	[NT]	LCS-5	72%
Toluene	mg/kg	0.5	GC.16	<0.5	[NT]	[NT]	LCS-5	81%
Ethylbenzene	mg/kg	1	GC.16	<1.0	[NT]	[NT]	LCS-5	88%
m+p-xylene	mg/kg	2	GC.16	<2.0	[NT]	[NT]	LCS-5	97%
o-Xylene	mg/kg	1	GC.16	<1.0	[NT]	[NT]	LCS-5	98%
Surrogate aaa-Trifluorotoluene	%		GC.16	72	[NT]	[NT]	LCS-5	114%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
sTPH in Soil (C10-C36)						Base II Duplicate II %RPD		
Date extracted	-			9/8/201 0	[NT]	[NT]	LCS-6	9/8/2010
Date analysed	-			9/8/201 0	[NT]	[NT]	LCS-6	9/8/2010
TPH C10 - C14	mg/kg	50	GC.3	<50	[NT]	[NT]	LCS-6	80%
TPH C15 - C28	mg/kg	100	GC.3	<100	[NT]	[NT]	LCS-6	88%
TPH C29 - C36	mg/kg	100	GC.3	<100	[NT]	[NT]	LCS-6	85%
Surrogate o-Terphenyl	%		GC.3	94	[NT]	[NT]	LCS-6	105%



QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
PAHs in Soil						Base II Duplicate II %RPD		j
Date extracted	-			9/8/201 0	[NT]	[NT]	LCS-6	9/8/2010
Date analysed	-			10/8/20 10	[NT]	[NT]	LCS-6	10/8/2010
Naphthalene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-6	102%
Acenaphthylene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Acenaphthene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Fluorene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-6	94%
Phenanthrene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-6	103%
Anthracene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Fluoranthene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-6	94%
Pyrene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-6	96%
Benzo(a)anthracene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Chrysene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	LCS-6	105%
Benzo(b+k)fluoranthene	mg/kg	0.2	GC.12 subset	<0.2	[NT]	[NT]	[NR]	[NR]
Benzo(a)pyrene	mg/kg	0.05	GC.12 subset	<0.05	[NT]	[NT]	LCS-6	111%
Indeno(1,2,3-c,d)pyrene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Dibenzo(a,h)anthracene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Benzo(g,h,i)perylene	mg/kg	0.1	GC.12 subset	<0.1	[NT]	[NT]	[NR]	[NR]
Surrogate p-Terphenyl-d ₁₄	%		GC.12 subset	76	[NT]	[NT]	LCS-6	126%

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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Speciated Phenols in Soil						Base II Duplicate II %RPD		
Date extracted	-			10/08/2 010	44271-1	10/08/2010 10/08/2010	LCS-6	10/08/2010
Date analysed	-			11/08/2 010	44271-1	11/08/2010 11/08/2010	LCS-6	11/08/2010
Phenol	mg/kg	1	GC.12	<1.0	44271-1	<1.0 <1.0	LCS-6	98%
2-Chlorophenol	mg/kg	1	GC.12	<1.0	44271-1	<1.0 <1.0	LCS-6	95%
2-Methylphenol	mg/kg	1	GC.12	<1.0	44271-1	<1.0 <1.0	[NR]	[NR]
3/4-Methylphenol	mg/kg	2	GC.12	<2.0	44271-1	<2.0 <2.0	[NR]	[NR]
2-Nitrophenol	mg/kg	1	GC.12	<1.0	44271-1	<1.0 <1.0	[NR]	[NR]
2,4-Dimethylphenol	mg/kg	1	GC.12	<1.0	44271-1	<1.0 <1.0	[NR]	[NR]
2,4-Dichlorophenol	mg/kg	1	GC.12	<1.0	44271-1	<1.0 <1.0	[NR]	[NR]
2,6-dichlorophenol	mg/kg	1	GC.12	<1.0	44271-1	<1.0 <1.0	[NR]	[NR]
2,4,5-trichlorophenol	mg/kg	1	GC.12	<1.0	44271-1	<1.0 <1.0	[NR]	[NR]
2,4,6-trichlorophenol	mg/kg	1	GC.12	<1.0	44271-1	<1.0 <1.0	[NR]	[NR]
2,4-dinitrophenol	mg/kg	10	GC.12	<10	44271-1	<10 <10	[NR]	[NR]
4-nitrophenol	mg/kg	10	GC.12	<10	44271-1	<10 <10	LCS-6	72%
2,3,4,6-tetrachlorophenol	mg/kg	1	GC.12	<1.0	44271-1	<1.0 <1.0	[NR]	[NR]
2-methyl-4,6-dinitrophen ol	mg/kg	10	GC.12	<10	44271-1	<10 <10	[NR]	[NR]
pentachlorophenol	mg/kg	10	GC.12	<10	44271-1	<10 <10	[NR]	[NR]
Surrogate 2-fluorophenol	%		GC.12	96	44271-1	64 68 RPD: 6	LCS-6	65%
Surrogate Phenol-d6	%		GC.12	96	44271-1	77 79 RPD: 3	LCS-6	72%
Surrogate 2,4,6-Tribromophenol	%		GC.12	60	44271-1	# #	LCS-6	46%
Surrogate p-Terphenyl-d ₁₄	%		GC.12	91	44271-1	107 103 RPD: 4	LCS-6	84%

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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Acid Extractable metals in soil						Base II Duplicate II %RPD		-
Date digested	-			09/08/2 010	[NT]	[NT]	LCS-2	09/08/2010
Date analysed	-			10/08/2 010	[NT]	[NT]	LCS-2	10/08/2010
Arsenic	mg/kg	4	Metals.20 ICP-AES	<4	[NT]	[NT]	LCS-2	102%
Cadmium	mg/kg	0.5	Metals.20 ICP-AES	<0.5	[NT]	[NT]	LCS-2	104%
Chromium	mg/kg	1	Metals.20 ICP-AES	<1	[NT]	[NT]	LCS-2	105%
Lead	mg/kg	1	Metals.20 ICP-AES	<1	[NT]	[NT]	LCS-2	103%
Mercury	mg/kg	0.1	Metals.21 CV-AAS	<0.1	[NT]	[NT]	LCS-2	99%
Nickel	mg/kg	1	Metals.20 ICP-AES	<1	[NT]	[NT]	LCS-2	105%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank
Moisture				
Date prepared	-			9/8/201 0
Date analysed	-			9/8/201 0
Moisture	%	0.1	LAB.8	<0.10

QUALITY CONTROL VOCs in Zero Headspace ASLP	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results Base II Duplicate II %RPD	Spike Sm#	Spike % Recovery
Date extracted	-			10/8/20 10	[NT]	[NT]	LCS-W1	10/8/2010
Date analysed	-			10/8/20 10	[NT]	[NT]	LCS-W1	10/8/2010
Styrene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	99%
Surrogate Dibromofluoromethane	%		GC.13	96	[NT]	[NT]	LCS-W1	92%
Surrogate toluene-d8	%		GC.13	93	[NT]	[NT]	LCS-W1	102%
Surrogate 4-BFB	%		GC.13	79	[NT]	[NT]	LCS-W1	101%

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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
BTEX in Zero Headspace						Base II Duplicate II %RPD		
Date extracted	-			10/8/20 10	[NT]	[NT]	LCS-W1	10/8/2010
Date analysed	-			10/8/20 10	[NT]	[NT]	LCS-W1	10/8/2010
Benzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	86%
Toluene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	100%
Ethylbenzene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	106%
m+p-xylene	µg/L	2	GC.13	<2.0	[NT]	[NT]	LCS-W1	110%
o-xylene	µg/L	1	GC.13	<1.0	[NT]	[NT]	LCS-W1	118%
Surrogate Dibromofluoromethane	%		GC.13	96	[NT]	[NT]	LCS-W1	86%
Surrogate toluene-d8	%		GC.13	93	[NT]	[NT]	LCS-W1	101%
Surrogate 4-BFB	%		GC.13	79	[NT]	[NT]	LCS-W1	106%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Metals-ASLP Neutral (ICP-MS)						Base II Duplicate II %RPD		
Date extracted	-			11/08/2 010	44271-1	11/08/2010 11/08/2010	LCS-W1	11/08/2010
Date analysed	-			11/08/2 010	44271-1	11/08/2010 11/08/2010	LCS-W1	11/08/2010
Arsenic in ASLP	µg/L	1	Metals.22 ICP-MS	<1	44271-1	8 8 RPD: 0	LCS-W1	106%
Cadmium in ASLP	µg/L	0.1	Metals.22 ICP-MS	<0.1	44271-1	<0.1 <0.1	LCS-W1	108%
Chromium in ASLP	µg/L	1	Metals.22 ICP-MS	<1	44271-1	7 7 RPD: 0	LCS-W1	88%
Lead in ASLP	µg/L	1	Metals.22 ICP-MS	<1	44271-1	<1 <1	LCS-W1	117%
Mercury in ASLP	µg/L	0.5	Metals.21 ASLP	<0.50	44271-1	<0.50 <0.50	LCS-W1	116%
Nickel in ASLP	µg/L	1	Metals.22 ICP-MS	<1	44271-1	2 2 RPD: 0	LCS-W1	86%



QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
sTPH in water leach						Base II Duplicate II %RPD		
Date extracted	-			5/8/201 0	[NT]	[NT]	LCS-W1	5/8/2010
Date analysed	-			5/8/201 0	[NT]	[NT]	LCS-W1	5/8/2010
TPH C10 - C14	µg/L	50	GC.3	<50	[NT]	[NT]	LCS-W1	83%
TPH C15 - C28	µg/L	100	GC.3	<100	[NT]	[NT]	LCS-W1	117%
TPH C29 - C36	µg/L	100	GC.3	<100	[NT]	[NT]	LCS-W1	92%
<i>Surrogate</i> o-Terphenyl	%		GC.3	128	[NT]	[NT]	LCS-W1	117%

QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
PAHs in water leach						Base II Duplicate II %RPD		
Date extracted	-			05/08/2 010	[NT]	[NT]	LCS-W1	05/08/2010
Date analysed	-			06/08/2 010	[NT]	[NT]	LCS-W1	06/08/2010
Naphthalene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	94%
Acenaphthylene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Acenaphthene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Fluorene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	109%
Phenanthrene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	101%
Anthracene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Fluoranthene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	96%
Pyrene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	106%
Benzo(a)anthracene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Chrysene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	104%
Benzo(b+k)fluoranthene in ASLP	mg/L	0.002	GC.12 ASLP	<0.002	[NT]	[NT]	[NR]	[NR]
Benzo(a)pyrene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	LCS-W1	113%
Indeno(1,2,3-c,d)pyrene - ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Dibenzo(a,h)anthracene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Benzo(g,h,i)perylene in ASLP	mg/L	0.001	GC.12 ASLP	<0.001	[NT]	[NT]	[NR]	[NR]
Surrogate p-Terphenyl-d14	%		GC.12	116	[NT]	[NT]	LCS-W1	99%

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QUALITY CONTROL	UNITS	PQL	METHOD	Blank	Duplicate Sm#	Duplicate results	Spike Sm#	Spike % Recovery
Speciated Phenols in water						Base II Duplicate II %RPD		
Date extracted	-			05/08/2 010	[NT]	[NT]	LCS-W1	05/08/2010
Date analysed	-			05/08/2 010	[NT]	[NT]	LCS-W1	05/08/2010
Phenol	µg/L	10	GC.12	<10	[NT]	[NT]	LCS-W1	34%
2-Chlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	LCS-W1	94%
2-Methylphenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
3/4-Methylphenol	µg/L	20	GC.12	<20	[NT]	[NT]	[NR]	[NR]
2-Nitrophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4-Dimethylphenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4-Dichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,6-Dichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4,5-Trichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4,6-Trichlorophenol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2,4-Dinitrophenol	µg/L	100	GC.12	<100	[NT]	[NT]	[NR]	[NR]
4-Nitrophenol	µg/L	100	GC.12	<100	[NT]	[NT]	LCS-W1	54%
2,3,4,6-Tetrachlorophen ol	µg/L	10	GC.12	<10	[NT]	[NT]	[NR]	[NR]
2-methyl-4,6-dinitrophen ol	µg/L	100	GC.12	<100	[NT]	[NT]	[NR]	[NR]
Pentachlorophenol	µg/L	100	GC.12	<100	[NT]	[NT]	[NR]	[NR]
Surrogate 2-fluorophenol	%		GC.12	66	[NT]	[NT]	LCS-W1	66%
Surrogate Phenol-d6	%		GC.12	32	[NT]	[NT]	LCS-W1	36%
Surrogate 2,4,6-Tribromophenol	%		GC.12	112	[NT]	[NT]	LCS-W1	116%
Surrogate p-Terphenyl-d ₁₄	%		GC.12	126	[NT]	[NT]	LCS-W1	119%
QUALITY CONTROL Speciated Phenols in Soil	UNIT	S	Dup. Sm#		Duplicate Duplicate + %RPI	Spike Sm#	Spike % Recovery	
Date extracted	-		[NT]		[NT]	44271-2	10/08/2010	=
Date analysed	_		[NT]		[NT]	44271-2	11/08/2010	
Phenol	mg/k	a	[NT]		[NT]	44271-2	75%	
2-Chlorophenol						44271-2	#	
-	mg/k		[NT]		[NT]			
2-Methylphenol	mg/k		[NT]		[NT]	[NR]	[NR]	
3/4-Methylphenol	mg/k		[NT]		[NT]	[NR]	[NR]	
2-Nitrophenol	mg/k	g	[NT]		[NT]	[NR]	[NR]	
2,4-Dimethylphenol	mg/k	g	[NT]		[NT]	[NR]	[NR]	
2,4-Dichlorophenol	mg/k	g	[NT]		[NT]	[NR]	[NR]	
2,6-dichlorophenol	mg/k	g	[NT]		[NT]	[NR]	[NR]	
2,4,5-trichlorophenol	mg/k	g	[NT]		[NT]	[NR]	[NR]	
2,4,6-trichlorophenol	mg/k	a	[NT]		[NT]	[NR]	[NR]	

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QUALITY CONTROL	UNITS	Dup. Sm#	Duplicate	Spike Sm#	Spike % Recovery
Speciated Phenols in Soil			Base + Duplicate + %RPD		
2,4-dinitrophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
4-nitrophenol	mg/kg	[NT]	[NT]	44271-2	#
2,3,4,6-tetrachlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
2-methyl-4,6-dinitrophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
pentachlorophenol	mg/kg	[NT]	[NT]	[NR]	[NR]
Surrogate 2-fluorophenol	%	[NT]	[NT]	44271-2	#
Surrogate Phenol-d6	%	[NT]	[NT]	44271-2	64%
Surrogate 2,4,6-Tribromophenol	%	[NT]	[NT]	44271-2	#
Surrogate p-Terphenyl-d14	%	[NT]	[NT]	44271-2	108%



Report Comments:

Total Petroleum Hydrocarbons in tclp: # Percent recovery is not possible to report as the high concentration of analytes in the sample/s have caused interference.

Total Petroleum Hydrocarbons in soil:# Percent recovery is not possible to report as the high concentration of analytes in the sample/s have caused interference.

Phenol's in soil by GCMS: # Percent recovery is not possible to report due to interference from analytes (other than those being tested) in the sample/s. Asbestos was analysed by Approved Identifier: Not applicable for this job INS: Insufficient sample for this test NT: Not tested PQL: Practical Quantitation Limit NR: Not requested <: Less than >: Greater than NA: Test not required

Quality Control Definitions

LCS: Laboratory Control Sample **RPD: Relative Percent Difference**

Blank: This is the component of the analytical signal which is not derived from the sample but from reagents, glassware etc, can be determined by processing solvents and reagents in exactly the same manner as for samples. Duplicate: This is the complete duplicate analysis of a sample from the process batch. If possible, the sample selected should be one where the analyte concentration is easily measurable.

Matrix Spike: A portion of the sample is spiked with a known concentration of target analyte. The purpose of the matrix spike is to monitor the performance of the analytical method used and to determine whether matrix interferences exist. LCS (Laboratory Control Sample): This comprises either a standard reference material or a control matrix (such as a blank sand or water) fortified with analytes representative of the analyte class. It is simply a check sample.

Surrogate Spike: Surrogates are known additions to each sample, blank, matrix spike and LCS in a batch, of compounds which are similar to the analyte of interest, however are not expected to be found in real samples.

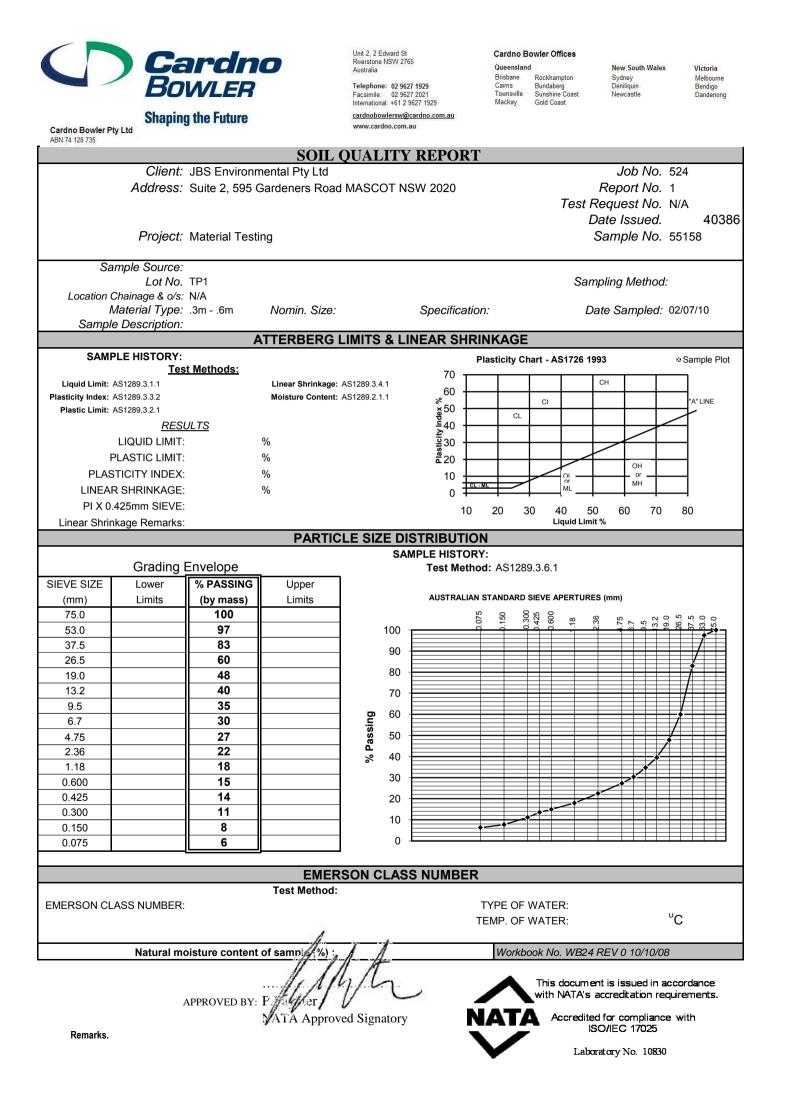
Laboratory Acceptance Criteria:

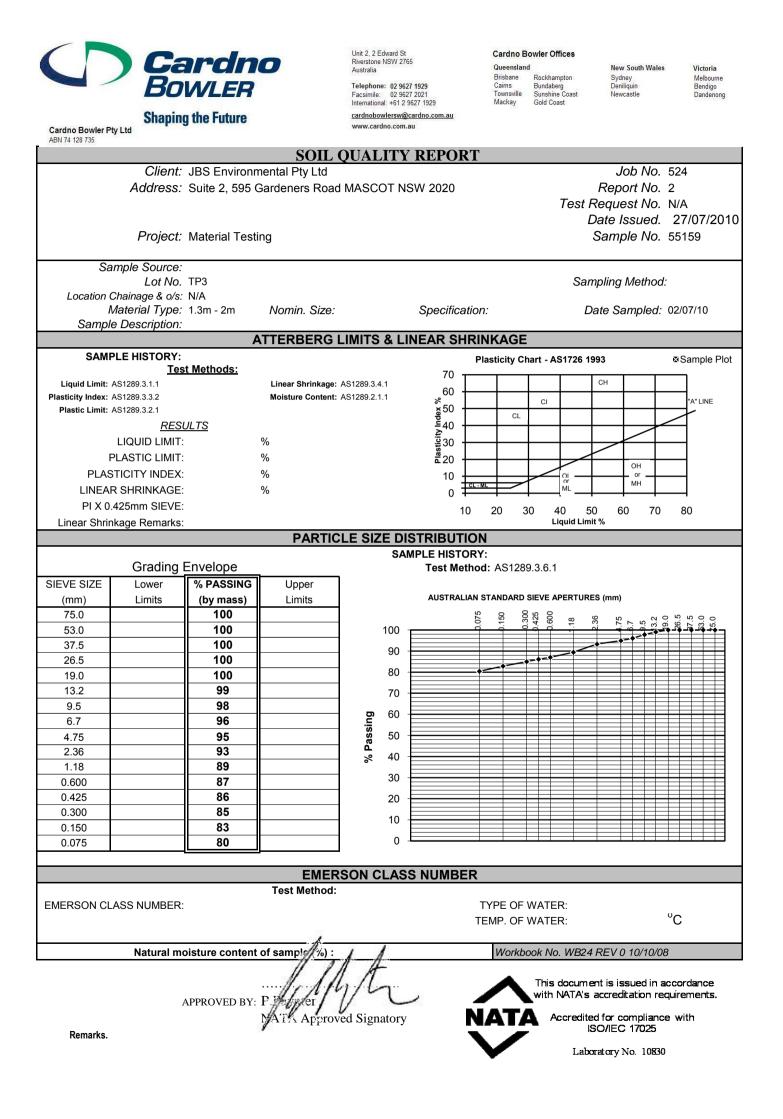
Duplicates: <5xPQL - any RPD is acceptable >5xPQL - 0-50% RPD is acceptable.

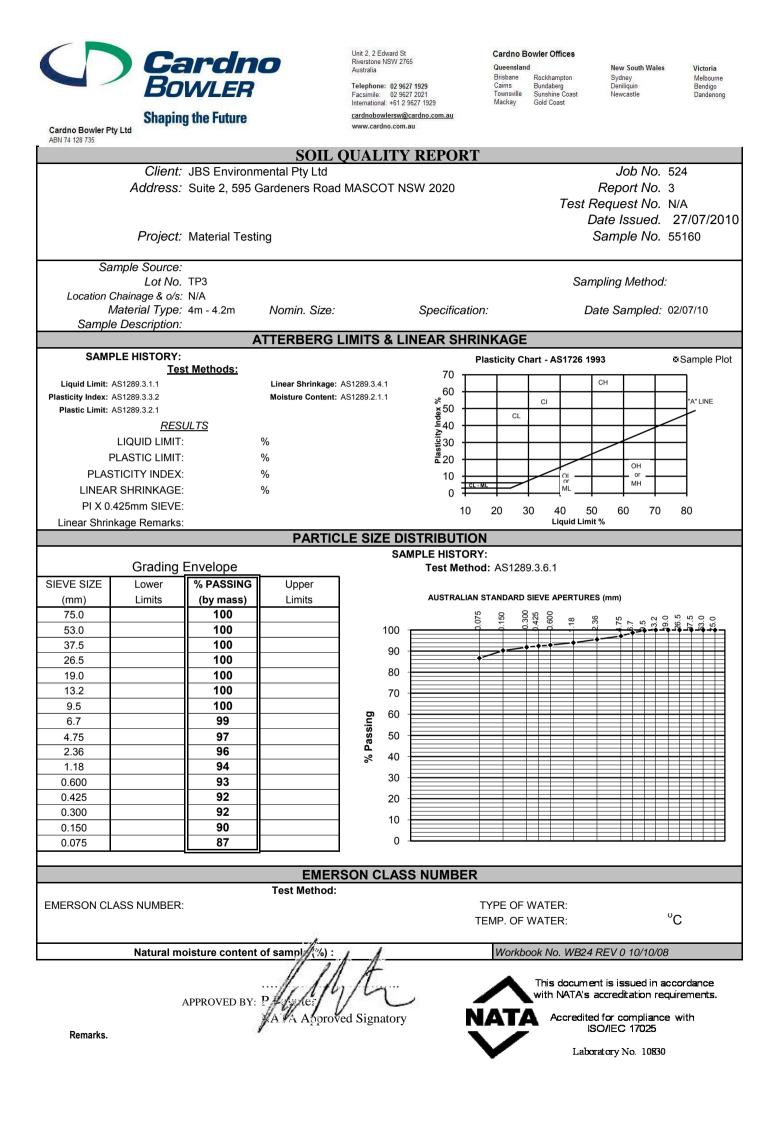
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Document Status

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No.		Name	Name	Signature	Date			
6	Sumi Dorairaj	Matthew Bennett	Matthew Bennett		24/06/11			
7	Sumi Dorairaj	Matthew Bennett	Matthew Bennett	Appentit	16/07/11			
8	Sumi Dorairaj	Matthew Bennett	Matthew Bennett	Appentet	16/07/11			



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