

Additional Ground Investigation for Land at Bankside, Banbury

Final Report

Prepared by

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for

Bovis Homes, Barratt Homes and Taylor Wimpey Homes

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EXECUTIVE SUMMARY AND CONCEPTUAL SITE MODEL

	Purpose of this	To review previous works at the site, add new data and identify any geo-environmental or
SITE INFORMATION & SETTING	report	geotechnical constraints associated with the site.
	Client	Bovis Homes, Barratt Homes and Taylor Wimpey Homes.
	Site	Land at Bankside, Banbury.
	Site Location	Land north east of Oxford Road, west of the Oxford Canal and east of Bankside in Banbury. Nearest postcode: OX15 4AD). Approximate National Grid Reference of centroid is 446470E 238326N.
'E INFOI	Current Land Use & Description	The site covers approximately 75ha and is currently agricultural land with a fenced off area of over grown open ground predominantly used by dog walkers at the north end of the site.
SIT	Development	Residential development in the south and north of the site with public open space on the central east part of the site.
	Site History	The area of land at the northern end of the site is known to formerly have been used as a tip. The remainder of the site is undeveloped farm land.
(yat	Geology	The southern end of the site is underlain by a thin layer of the Whitby Mudstone Formation and the Marlstone Rock Bed. The central part of the site is underlain by the Dyrham Formation. The north and eastern part of the site is underlain by the Charmouth Mudstone Formation. There is Made Ground associated with the former tip in the northern end of the site.
PHASE 1 (DESK STUDY)	Hydrogeology	The Made Ground is considered to be low or low to moderate permeability because of high clay content. The Whitby Mudstone Formation is classified as Unproductive Strata. The Marlstone Rock Formation is classified as a Secondary A Aquifer. The Dyrham Formation is classified as a Undifferentiated Secondary Aquifer. The Charmouth Mudstone Formation is classified as Unproductive Strata. The site is not in a source protection zone. Springs issue from of the Marlstone Rock Bed / Dyrham Formation junction with the Charmouth Mudstone.
_	Hydrology	The nearest watercourse is the Oxford Canal located adjacent to the north eastern boundary of the site.
	Environmental Sensitivity	The Oxford Canal and the land beyond the Oxford Canal are registered as an Environmentally Sensitive Area.
	Historical Site Works	Historical assessments of the site have been undertaken by Wardell Armstrong LLP and Corsair since 2005. The data from theses assessments are used alongside Hydrock data in this report.
TION	Hydrock Site Works	The Hydrock Ground investigation comprised 41 trial pits, with geo-environmental chemical testing, excavated to a maximum depth of 3.5m bgl.
PHASE 2 – GROUND INVESTIGATION	Findings of the Ground Investigation	In the Tip Area Made Ground was encountered over the Charmouth Mudstone Formation. In general, the Made Ground consisted of variable content comprising gravelly clay with cobbles and boulders, gravel and cobbles and boulders of brick, white polystyrene, plastic sheeting, concrete, concrete fragments, plastic and a boulder-sized section of brick walling. Elevated concentrations of metals and PAHs were recorded within the Made Ground materials. The elevated metals are considered to the attributed to the natural background levels in the local area. The Charmouth Mudstone Formation comprised variable conditions including low to high strength, orange brown, brown and grey, silty, residual clay and sand with high groundwater seepage. The Dyrham Formation comprised very low to low strength, orange brown and grey, silty, residual clay overlaying the Marlstone Rock Bed. The Whitby Mudstone Formation comprised medium to high strength, firm to stiff, brown and yellow brown mottled, gravelly, sandy, silty, residual clay overlaying the Marlstone Rock Bed. The Marlstone Rock Bed comprised medium to high strength, red brown and brown, sandy, gravelly clay and moderately strong, blue grey and red brown banded, shelly, weathered limestone. Elevated concentrations of vanadium and arsenic were recorded within the natural soils however these were found not to be bioaccessible and are attributed to the natural background levels of the site.

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	Site Preparation, Earthworks &	Where the proposed development is in close proximity to existing live services, they must be located and a way leave clearly marked prior to undertaking ground works in the vicinity.
	Landscaping	Temporary slope stability works may be required in any deep excavations in the north of the site.
		Obstructions may be present in the Made Ground .
		Significant excavation into the Marlstone Rock Bed may require the use of pneumatic breakers or rock rippers and there will be a reduction in progress where their use is necessary.
		Topsoil and unsuitable Made Ground should be removed from beneath all building and hard standing areas.
		Spoil resulting from excavations within the Made Ground may be suitable for reuse as fill to raise site levels elsewhere within the tip area, subject to its suitability with respect to appropriate geotechnical and geo-environmental specifications. Contaminated material should not be re-used on any part of the site currently uncontaminated.
		Well-pumping may be required for dewatering as sump pumping will increase the instability of unsupported ground in these conditions.
		It is recommended that no site personnel enter any trenches unless there is adequate support and this has been assessed by a competent person.
		The earthworks will need to be undertaken under a Materials Management Plan to ensure the appropriate management and reuse of the existing soils. Where the earthworks perform a structural role, a suitable specification will be required.
	Foundations	Tip area - it is anticipated that trench fill spread foundations should be appropriate for the proposed row of plots along the western boundary of the tip area, to be founded in the underlying natural soils with the remainder of the tip area requiring deeper pile foundations to penetrate through the Made Ground and the unstable ground conditions in the eastern end of the tip area into the stiff clay of the Charmouth Mudstone.
ATIONS		Northern area outside of the tip - competent founding strata ranges from 0.2 to 3m bgl with no obvious pattern. As a result of the variable ground conditions a combination of spread foundations and piled foundations would be appropriate in this area. This would need to be assessed by a plot by plot basis.
NSIDER		Southern area - the underlying Marlstone Rock Bed should be adequate to support the proposed development using spread foundations.
ASSESSMENT & CONSIDERATIONS	Floor Slabs	In the north part of the site as Made Ground is greater than 600mm thick in the tip area and clay soils of moderate to high volume change potential are present, it is recommended that suspended floor slabs should be adopted, in accordance with NHBC Standards. Ground bearing slabs may be used in the south of the site if, the foundation depth (such as due to the influence of trees) is less than 1.5m, any fill is suitable, well-compacted granular material and less than 600mm thick and it is demonstrated that desiccation is not present and soils are at their equilibrium moisture content.
ASS	Roads	In the north part of the site where the CBR was found to be less than 2.5%, the sub-grade may be unsuitable for both the trafficking of site plant and as support for a permanent foundation, without improvement works being undertaken. Improvement works should be carried out.
		It is considered likely a CBR of 5% will be achievable over the south of the site of the site and can be used for preliminary design, subject to in situ testing during construction. Proof rolling of the formation level will be required and any loose or soft spots to be removed and replaced with an engineered fill, in accordance with a suitable specification.
	Soakaways & Drainage	The 2012 Wardell Armstrong highlights that limited soakaway testing was undertaken at the site but details are not available and it is possible that further soakaway testing may be required at the site.
	Buried Concrete	The materials at the site would be classified as Design Sulfate Class DS-1 and Aggressive Chemical Environment for Concrete (ACEC) Class AC-1d.
	Results of Generic Risk Assessment	Human Health - PAHs and potential asbestos containing materials in the Made Ground associated with the tip area in the north of the site.
	Requiring Consideration	Elevated naturally occurring vanadium in the areas underlain by the Marlstone Rock Bed and Whitby Mudstone Formation are not considered to represent a risk to human health, but this has yet to be confirmed by the Local Authority.
		Controlled Waters - None.
		Plant Life - None.
		Human health / property from ground gases - Full radon protective measures are indicated by the current guidance.
	Outline Contamination	A cut and fill remedial strategy for the tip area will have to be developed in consultation with the design team and the regulatory authorities.
	Remediation	The material removed, if geotechnical suitable could be used to raise levels under roads and hard

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	Strategy	standing or with proposed public open space as long as a suitable cover or contaminant pathway breakage is in place to prevent direct contact with the Made Ground by end users of the site.
		Through an appropriate Materials Management Plan and on-site processing of materials the development will keep as much of the Made Ground material on site as possible, thus minimising vehicle movements and landfilling.
		The 'fine' fraction, containing the soil, will require sampling (e.g. one sample per 500m ³) to aid in the assessment of its suitability for reuse on the site as either clean cover or, if the material is contaminated, as fill material to raise the site levels, where required and be covered by an appropriate clean cover.
		Areas of Made Ground under gardens and public open space will require a clean cover of a minimum of 600mm thickness. The extent and thickness will need to be estimated from knowledge of the existing and final ground levels.
		Foundations will have to be taken down to natural ground and consideration given to protection of water supply pipes and other utilities.
		Full radon protection measures will be required and these should be capable of preventing ingress of naphthalene vapours in the tip area.
		The methodology for the remediation should be detailed in a Remediation Method Statement which will need to be submitted to the NHBC and the regulatory authorities for approval.
		Following completion of the above works verification reports, undertaken by a suitably qualified independent engineer will be required.
Î	Construction Materials & Water Supply Pipework	It is envisaged that standard pipework will be suitable for the majority of the site, but barrier pipe will be required within the Made Ground. However, due to conflicting guidance, confirmation should be sought from the water supply company at the earliest opportunity.
	Uncertainties	Risks to human health from naturally occurring vanadium is not considered significant, but at the time of writing this has yet to be confirmed with the Local Authority.
		Asbestos-containing materials may be present in the Made Ground.
		Risks to the surface water environment are not considered significant, but this will have to be agreed with the Environment Agency. A piling risk assessment is likely to be required in the tip area to demonstrate the lack of establishing any new contaminant pathway.
		There is the potential for running sand conditions to develop in excavations close to the canal where silty or sandy soils are combined with a high water table. In these conditions it may be necessary to employ well-point dewatering before stable excavations can be formed.
		The practical cut-off between trenchfill and piled foundations is normally taken to be a founding depth of 2-2.5 mbgl. It has been shown that potential foundation depths exceed this threshold in the northern part of the site. The areas where piling is likely to be required have been highlighted, but this is a provisional finding and more detailed assessment will be needed on a ploy by plot basis. Existing CBR testing is sufficient for preliminary design, but confirmatory testing will be required in
		actual road foundation locations.
		The 2012 Wardell Armstrong highlights that limited soakaway testing was undertaken at the site but details are not available and it is possible that further soakaway testing may be required at the site.
	Further Work	The following further works will be required with regard to the tip area in the north part of the site:
		 Liaison with the NHBC, Contaminated Land Officer and Environment Agency concerning the findings of this report in respect of contamination.
		Confirmation with the water company with respect to water supply pipe materials.
		Provision of a piling contamination risk assessment.
		• Provision of a Geotechnical Design Report including design and possible further geotechnical investigation for (a) spread footings (including suitability of plots, safe bearing pressure and mitigation of the effects of trees and shrubs); and (b) piled foundations (including specialist sub-contractor design and design of a piling mat for tracked plant).
		 Soakaway testing in areas where these may be feasible, ie granular soils without high water table.
		Provision of a Materials Management Plan for re-use of site arisings.
		 Provision of a Remedial Method Statement for the cover over the tip area and agreement with the regulatory authorities; and
		Verification of the remedial works on completion.
This Fr	an ting Common (and	l s part of Hydrock Consultants Limited report number R/12702/001 (Issue 2) and should not be used

This Executive Summary forms part of Hydrock Consultants Limited report number R/12702/001 (Issue 2) and should not be used as a separate document.

1.0 INTRODUCTION

1.1 <u>Terms of Reference</u>

In November 2012, Hydrock Consultants Limited (Hydrock) was commissioned by Bovis Homes, Barratt Homes and Taylor Wimpey Homes (E7556/002) to undertake additional ground investigation at Land at Bankside, Banbury. The site is located on land north east of Oxford Road, west of the Oxford Canal and east of Bankside in Banbury (nearest postcode : OX15 4AD).

The site covers approximately 75ha and is currently predominantly agricultural land with a fenced off area of overgrown open ground predominantly used by dog walkers at the north end of the site.

The proposed development is to comprise combined residential and commercial development with public open space.

A site location plan (Drawing C12702-G001), is presented in Appendix A.

1.2 Objectives

The objectives of this investigation are to review previous works at the site and to identify any geo-environmental or geotechnical constraints associated with the site.

1.3 <u>Scope</u>

The scope of work for this commission comprises:

- a review of historical works and a site walk over reconnaissance;
- a review of previous investigations carried out at the site;
- a supplementary ground investigation including trial pitting and chemical testing to provide confirmation of the findings of the historical investigations and provide additional information to support any remedial proposals; and
- reporting on findings of the review, ground investigation, geo-environmental assessment of the site conditions.

See Appendix D for detailed reporting methodology.

1.4 Information Sources Provided by the Client

A number of phases of site investigations have been undertaken at the site. In preparing this report Hydrock has consulted the following documentation:

- October 2005 site investigation by Corsair Consultants focused on the former tip area within the north of the site (reviewed as part of the WA 2010 Report);
- 2006 site investigation by Corsair Consultants was designed to provide a broader assessment of the ground conditions across the site (reviewed as part of the WA 2010 Report);

- 2007 site investigation and testing by Wardell Armstrong LLP, Further investigation works were then undertaken to establish the bio-accessibility of the arsenic present within the site (reviewed as part of the WA 2010 Report);
- August 2010, Wardell Armstrong LLP, Report on Ground Conditions, Bankside, Banbury;
- 3 February 2012, Wardell Armstrong Letter Report, Assessment of Area Adjacent to Fuel Station Bankside, Banbury;
- 26 March 2012, Wardell Armstrong Letter Report, *Site at Bankside, Banbury Results of Ground Gas Monitoring*;
- 17th May 2012, GEG Geo Environmental Group, *Intrusive Assessment of Canal Base*, Oxford Canal, Banbury, Oxfordshire; and
- August 2012, Wardell Armstrong LLP, *Remediation, Ground Investigation Report*, Bankside, Banbury;
- 13-09-2012, Focus, Drawing; SK01 Development Proposals Parcel E -A1L;
- 22-10-2012, Focus, Drawing; SK02 Development Proposals-A0L;
- 20-10-2012, Focus, Drawing; SK03 Development Proposals-A0L;
- 23-10-2012, Focus, Drawing; SK04 Development Proposals Parcel F-A0L;
- 23-10-2012, Focus, Drawing; SK05 Development Proposals Parcel D;
- 31-10-2012, M-EC, Drawing; 20488_00_001 Engineering Viability Plan Sheet 1 Of 2;
- 31-10-2012, M-EC, Drawing; 20488_00_002 Engineering Viability Plan Sheet 2 Of 2;
- July 2004, Gallagher Estates, Drawing; Bodicote, Banbury, Oxfordshire Constraints Plan;
- 29-06-2010, Paul Drew Design, Drawing; 005 College Fields, Bankside Banbury, Master Plan; and
- 25-09-2012, SAVILLS, Drawing; Longford Park, Banbury Extract.

This report covers all previous investigations at the site which are summarised in Section 2.8.

2.0 BACKGROUND

The available information from previous reports has been reviewed and is presented in this section, augmented by additional information from Hydrock's resources (Sections 2.1 to 2.5).

2.1 <u>Site Referencing</u>

The site is referenced in Table 2.1.

Item	Brief Description	
Site name	Land at Bankside, Banbury.	
Site location and Grid reference	The site is located 2km to the south of Banbury town centre to the west of the Oxford Canal. The site occupies an area of approximately 75 ha and is shown on Drawing C12702-G001. Nearest postcode is OX15 4AD. Approximate National Grid Reference of centroid is 446470E 238326N.	

A site location plan is provided in Appendix A (Drawing C12702-G001).

2.2 Site Description and Walk-Over Survey

The basic description of the current site conditions is summarised in Table 2.2.

Item	Brief Description	
Site access	Either via Canal Lane at the centre of the site or off of the road Bankside which follows the north western boundary of the site.	
Land area	Approximately 75 ha.	
Elevation, topography etc.	hy The elevation of the site ranges from approximately 123.5m AOD in the west the site to 89.7m AOD in the north east of the site. The southern half of the site is predominately flat. The north half of the site slopes downhill to the Oxfor Canal which is adjacent to the north eastern boundary of the site. Springs were located in east of the site which were noted to run down the site of the hedgerows towards the Oxford Canal.	
Site boundaries	Site boundaries consist of the Oxford Canal on the north eastern boundary the site with the remainder consisting of hedge boundaries either to roads (w of the site) or more agricultural fields to the south east of the site.	
Present land use	The site is currently open fields used primarily for agriculture.	
Surrounding land	To the west and south of the site is residential development, to the east of the site are agricultural fields and to the north east of the site is the Oxford Canal beyond which are more agricultural fields and the River Cherwell which lies approximately 230m beyond the canal.	

Table 2.2: Site Description

A walk-over reconnaissance survey was undertaken to confirm the findings of the information review and assess visually any potential hazards and receptors that may not have been picked up in the previous work. Photographs are presented in Appendix B.

2.3 <u>Geology</u>

The general geology of the site area is shown on the 1:50,000 geological map of Chipping Norton (Sheet 218) and is summarised in Table 2.3.

No superficial geology is indicated at the site and Drawing C12702-G004 (sourced from the previous site investigation of the site) illustrates the conjectured geological setting at the site.

Location	Age	Stratigraphic Name	Description
	Jurassic	Whitby Mudstone Formation - Mudstone.	The geological records indicate a small outlier of the Whitby Mudstone Formation in the south eastern part of the site.
	Jurassic	Marlstone Rock Formation Ferruginous Limestone And Ironstone	The Marlstone Rock Bed is shown to crop out in the southern half of the site. The Marlstone Rock Bed contains ironstone deposits. Weathered ironstone deposits are known to be associated with elevated levels of metals/metalloids, especially arsenic and vanadium in soils. Generally, areas which are rich in ironstone have naturally elevated levels of arsenic and vanadium in soils.
On site	Jurassic	Dyrham Formation- Siltstone And Mudstone, Interbedded.	There is an outcrop of the Dyrham Formation recorded as clays and silts, in the central portion of the site.
	Jurassic	Charmouth Mudstone Formation Mudstone.	There is an outcrop of the Charmouth Mudstone Formation in the northern portion of the site. It is reported that the junction between the Charmouth Mudstone Formation and the Dyrham Formation is marked by a change in slope and water seepages which are in places pronounced by a spring line. The geological map notes two springs (however, up to four springs are noted on some of the historic plans) within the site area near this junction.

Table 2.3: Geology

Due to the historic tipping in at the north end of the site, deep Made Ground is anticipated in this area. The depth of Made Ground is considered be up to 4m deep. Available information indicates that tipped materials included brick, concrete, timber, ironstone, tarmac, tile, asbestos, flint, ash, coal, slate and metal in a gravelly, sandy, clayey, matrix in this area.

2.4 BR 211 Guidance on Radon

Reference to the Annex A maps in BR 211 (Scivyer 2007), based on the Indicative Atlas of Radon in England and Wales (Miles *et a*l 2007) indicates that full radon protection is required for new dwellings at this location in line with current guidance.

The documentation indicates that between 10% and 30% of houses record radon levels above the action level of 200 bq/m^3 , therefore full radon protective measures would be required within the construction of residential dwellings at the Bankside site.

2.5 <u>Hydrogeology</u>

The strata beneath the site are classified by the Environment Agency as a Secondary Aquifer and /or Unproductive Strata, depending on the underlying geology. The aquifer designations given in Table 2.4 are based on the Environment Agency interactive aquifer designation map.

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Stratum	Aquifer Designation	Hydraulic Characteristics		
Made Ground (north end of the site)	N/A	Moderate to high porosity because of unconsolidated nature, but permeability likely to be constrained to low or low to moderate because of poor sorting and high clay content.		
Whitby Mudstone Formation - Mudstone.		The Whitby Mudstone Formation encountered in the area is mainly clay drift deposits with low permeability that has negligible significance for water supply or river base flow.		
Marlstone Rock Formation - Ferruginous Limestone And Ironstone	Secondary A Aquifer	The Marlstone Rock Bed is the likely source of the springs noted in the eastern end of the site which is the likely change in the geology to the Dyrham Formation and the Charmouth Mudstone Formation.		
Dyrham Formation - Siltstone And Mudstone, Interbedded.	Undifferentiated Secondary Aquifer	Secondary Undifferentiated has been assigned in cases where it has not been possible to attribute either category A or B to a rock type. The groundwater springs may be issuing at the boundary between the Marlstone Rock Formation and the Dyrham Formation or the Charmouth Mudstone Formation because of the variable nature of these strata.		
Charmouth Mudstone Formation - Mudstone.	Unproductive Strata	The Charmouth Mudstone Formation consists sedimentary bedrock deposited as mud, silt, sand and gravel rock layers with low permeability that have negligible significance for water supply or river base flow.		

Table 2.4: Hydraulic Characteristics of Strata

According to the Environment Agency web site, there are no recorded source protection zones within the site area.

Natural springs were noted in the eastern end of the site about halfway up the slope of the hill. It is likely that these springs issue from where the boundary of the Marlstone Rock Formation meets with either the Dyrham Formation of the Charmouth Mudstone Formation. It is therefore considered that groundwater within these Secondary Aquifers issue from the springs in the area.

Reference to the Environment Agency web site shows the following Groundwater Body beneath the site and the current chemical status.

Category	Label / Status
Waterbody ID	GB40602G600200
Waterbody name	Banbury Jurassic
River basin district	Thames
Current quantitative status	Good
Current chemical status	Good
Upward chemical trend	No
2015 predicted qualitative status	Good
2015 predicted chemical status	Good

 Table 2.5: Groundwater Body

Category	Label / Status
Overall risk	Probably Not At Risk
Protected area	Yes

The site is not within a Source Protection Zone (SPZ).

It is indicated that there are no licensed groundwater abstractions within 1 km of the site.

2.6 <u>Hydrology & Flooding</u>

The nearest watercourse shown on current plans is the Oxford Canal which forms the eastern boundary of the site. The River Cherwell lies approximately 230m beyond the Oxford Canal to the east of the site and is classified as Grade B by the Environment Agency. A number of springs (up to four) are identified to issue within the site. Additionally, anecdotal evidence from Cherwell District Council, reported in the Environmental Statement submitted with the Planning Application, suggests that minor ephemeral shallow ponds form from time to time in two stretches along the Oxford Canal.

Reference to the Environment Agency web site shows the following River Basin Management Plans – Rivers Current Ecological Quality information for the Oxford Canal adjacent to the east boundary of the site and the current chemical status (Table 2.6).

Category	Label / Status
Water Body Name	Oxford Canal, summit to Aynho
Waterbody Name	GB70610197
Management Catchment	Oxford Canal, summit to Aynho
River Basin District	N_A
Typology Description	Thames
Hydromorphological Status	Canal
Current Ecological Quality	Artificial
Current Chemical Quality	Good Potential
2015 Predicted Ecological Quality	Does Not Require Assessment
2015 Predicted Chemical Quality	Good Potential
Overall Risk	Does Not Require Assessment
Number of Measures Listed (waterbody level only)	-

Table 2.6: Surface Water Body

A significant (Category 2) pollution incident to Controlled Waters occurred approximately 188m to the east of the site boundary. The incident is not detailed fully within the report but described as involving 'chemicals'. The incident date is not supplied, however it is unlikely that this incident would affect the site. Further incidents have been recorded further from the site. It is unlikely that any of these occurrences would adversely affect the site.

A number of discharge consents are recorded within the area of the site with the closest being approximately 158m from the boundary of the site. The majority of these consents are

to water courses principally being the Oxford Canal and the River Cherwell. It is considered unlikely that these discharges would affect the site.

Several abstraction licences are in place within the area surrounding the site, the most significant of these is a potable water supply extracted from groundwater located approximately 1km from the site boundary. A further abstraction approximately 342m from the site is for the washing of vegetables.

The site is in Flood Zone 1, with low probability of flooding, however the site is located adjacent to both a Flood Zone 2, with medium/moderate probability of flooding and a Flood Zone 3, with high/significant probability of flooding located beyond the Oxford Canal to adjacent to the eastern site boundary.

No further consideration of flood risk is undertaken in this report. Specialist flood risk advice should be sought with regards to drainage and flooding.

2.7 Site History

The previous assessment of the site identified the area of land at the northern end of the site is known to formerly have been used as a tip. Available information indicates that tipped materials included brick, concrete, timber, ironstone, tarmac, tile, asbestos, flint, ash, coal, slate and metal in a gravelly, sandy, clayey matrix in this area. It is thought that the materials were tipped on to the natural surface and may be up to 4m thick in places. However, there is no official record of the materials that have been infilled in this area, nor is the site known to be a registered landfill site.

2.8 <u>Waste Management and Hazardous Substances</u>

A landfill is recorded approximately 400m to the north of the site boundary. The waste type is described to be domestic and inert materials and liquid sludge. It is not recorded when the landfill site was in operation nor whether the site has closed. It is considered unlikely that this landfill site would affect the Bankside site.

An existing petrol station is noted within the Envirocheck Report in the Wardell Armstrong 2010 report as being situated adjacent to the south western boundary of the site on the Oxford Road.

The land beyond the Oxford Canal is part of an Environmentally Sensitive Area Agreement and is a registered Environmentally Sensitive Area.

2.9 Previous Site Investigations or Other Reports

As outlined in Section 1.4 there has been significant previous ground investigation and assessment undertaken at the site. This is summarised below.

2.9.1 October 2005 Site Investigation by Corsair Consultants

In October 2005, Corsair Consultants focussed on the former tip area within the north of the Bankside site. These works comprised the excavation of 18 trial pits using a JCB 3CX excavator. The trial pits encountered generally Made Ground materials which comprised

gravelly and cobbly, sandy clay materials with varying proportions of broken brick, concrete and other construction debris. Geochemical testing of materials recovered from the investigation locations was undertaken to provide information on the contaminative nature of the materials.

In addition to the geochemical testing a spiker bar gas survey was undertaken at the location of each trial pit to a depth of 1m.

The trial pits identify multifarious anthropogenic materials in the tip area of the site up to 4m in thickness including brick, concrete, timber, ironstone, tarmac, tile, asbestos, flint, ash, coal, slate and metal in a gravelly sandy clayey matrix. The spiker bar gas survey revealed only low concentrations (up to 1.7%) of carbon dioxide and no methane. A total of fourteen trial pits encountered tipped materials across an area of approximately 3.3ha.

Decaying vegetation was found at the base of the tipped materials in a number of locations implying that the materials were deposited on the natural surface of the site, thus raising levels in this area of the site.

Geochemical testing results revealed elevated levels of arsenic, lead, PAH and TPH in selected samples compared to conservative screening values (i.e. SGV/GAC derived for residential with plant uptake standard land uses). Elevated levels of arsenic were detected in both Made Ground and natural soils at the site. Five samples were also analysed for asbestos fibres where asbestos containing material was suspected. Of the five samples tested, four returned positive results for asbestos fibres within cement bound materials.

In addition, ten soil samples were subjected to leachate testing for a range of inorganic and organic determinands. While organic analysis is normally not considered appropriate due to the processes involved in producing the leachate, as analysis had been undertake, results were included for initial assessment purposes. The leachate testing results were compared with water quality guideline values. None of the leachate results were above the relevant guidelines indicating that the soils in the tip area are not a significant risk to controlled waters.

2.9.2 2006 Site Investigation by Corsair Consultants

In April 2006 further site investigation designed to provide a broader assessment of the ground conditions across the site was undertaken by Corsair Consultants. The investigation involved the drilling of four cable percussion boreholes, excavation of forty two trial pits, undertaking five California Bearing Ratio (CBR) tests and four soakaway tests. The investigation works are reviewed in greater detail within the Wardell Armstrong 2010 'Report on Ground Conditions' with the testing results and logs included within that report.

The exploratory holes in this phase of investigation record varying geologic conditions across the site area broadly consistent with the published geology. All locations record natural material below the top soil across the site area investigated.

Soil, leachate, groundwater and gas samples were collected during the site investigation works for geochemical and geotechnical (soil only) analysis. A similar suite of chemical analysis was used as in the October 2005 investigation on selected samples.

Arsenic was generally found to be elevated in natural materials across the site, with elevated nickel in one sample. The recorded concentrations of arsenic (and nickel) increase in the southern site area; and are highest in soils above Marlstone Rock Bed. The arsenic (and nickel) was considered to be naturally occurring and to be associated with the weathering of the Marlstone Rock Bed (e.g. ironstone deposits) known to contain elevated concentrations of these substances.

The leachate testing results from this phase of investigation was compared to the water quality guidelines for screening purposes. As with the northern site area, all leachate samples record concentrations of all determinands were below the relevant water quality guidance.

Geotechnical testing on selected samples included natural moisture content, Atterberg limits, particle size distribution, particle density, dry density/moisture content relationship and oedometer consolidation testing.

2.9.3 2007 Site Investigation and Testing by Wardell Armstrong LLP

Further investigation works were undertaken to establish the bio-accessibility of the arsenic present within the site (reviewed as part of the WA 2010 Report).

In order to assess the potential significance of elevated arsenic recorded at the site further sampling was undertaken in the southern site area. Soil samples were collected from areas of the site where arsenic was known to be elevated. The soil samples were subject to geochemical analysis for total and PBET for arsenic.

The results enabled the generic assessment criteria to be modified to produce a Site Specific Assessment Criteria (SSAC). Through the development of Site Specific Assessment Criteria (SSAC) which took into account the PBET results it was demonstrated that the elevated concentrations of arsenic were within the SSAC.

This work demonstrated that the naturally occurring arsenic levels encountered at the site do not pose an unacceptable risk to human health.

2.9.4 August 2010, Wardell Armstrong LLP, Report on Ground Conditions, Bankside, Banbury

This report presents relevant background information regarding the geo-environmental setting of the site in the form of desk study researches and a review of published environmental information including hydrology, hydrogeology, radon, geology along with a site history and the findings from all the previous site investigations.

The findings from all the previous site investigation data at the site are used along with the understanding of the site and its environs from the desk study work to develop a Conceptual Site Model (CSM) and to design a remediation strategy with respect to the proposals for redevelopment.

The results of the risk assessments and revised understanding of the conceptual site model indicate that the levels of contaminants recorded in the former tip area Made Ground

materials are considered unsuitable for the residential development proposed for that area of the site without mitigation.

The risk assessment also demonstrated that elevated arsenic, considered to be naturally occurring, in the southern site area does not represent a significant risk to human health at the site and therefore no mitigation is required.

The report also recommends that it is necessary to provide full radon protection measures at the site in accordance with guidance from the BRE and highlights the potential risk from the fuel station adjacent to the western site boundary along with the potential for gas generation from the former tip in the northern part of the site.

An outline remediation strategy has been presented to mitigate the potential contamination issues in the north part of the site. The proposed remediation strategy includes:

- screening of material to remove deleterious materials from the soils, which would then be removed from site, if necessary;
- concrete and brick recovered from the screening process to be crushed and reused within the development of the site; and
- the 'fine' fraction, containing the soil, would require sampling (e.g. one sample per 500m³) to aid in the assessment of its suitability for re-use on the site.

The report concludes that depending on the results of the testing, it is likely that the majority of the soils generated from the screening process would be suitable for reuse in either residential, commercial or public open space areas of the site, either at depth or in the near surface, allowing the majority of materials to be retained in a sustainable approach.

Finally the report concludes that where necessary, depending on the extent of the excavations and subsequent processing of materials in the northern site area, a clean cover pathway break may be required to separate end users (this area of the site is proposed for residential development) from the underlying materials, if tipped materials are left in place. The extent of areas requiring clean cover (if any) would need to be developed following delineation of this area along with assessment of current and proposed finished ground levels in the development.

2.9.5 3 February 2012, Wardell Armstrong Letter Report, Assessment of Area Adjacent to Fuel Station – Bankside, Banbury

Additional site investigation works were undertaken by Wardell Armstrong within the vicinity of the fuel filling station and the car dealership located adjacent to the western boundary of the Bankside site.

The investigative works were aimed at determining whether this area of the site might have been affected by possible leaking petroleum hydrocarbons.

The site investigation works comprised the excavation of seven machine excavated trial pits to a maximum depth of 2.8m below ground level along the line of the boundary between the site and the filling station.

The trial pit excavations encountered natural soil and rock materials comprising red brown, sandy gravel and cobbles of sandstone and limestone in a clay matrix underlain by weathered rock recovered as gravel and cobbles of limestone and sandstone. No visual or olfactory evidence of hydrocarbons were noted within any of the trial pits at the time of the site investigation.

A total of twelve samples of soils from the seven trial pit excavations were scheduled for a suite of laboratory geochemical testing. The results indicate that none of the determinands were found above the laboratory limit of detection. This confirms that petroleum hydrocarbon compounds are not present at detectable levels within the soils adjacent to the boundary of the site with the filling station.

The report concluded that the results of the investigations provide sufficient evidence that the site is not being significantly impacted by fuel and or fuel derived compounds emanating from the filling station.

2.9.6 26 March 2012, Wardell Armstrong Letter Report, Site at Bankside, Banbury – *Results of Ground Gas Monitoring*

This ground gas monitoring letter report presents the results of the ground investigation to assess ground gas generation associated with the tip area in the northern part of the site.

The purpose of the report was the assessment of the current near-surface soil gas regime in the tip area with respect to the proposed future residential development.

The site investigation included the installation of 20 boreholes on a 50m grid across the former tip area of the site to depths ranging between 3 and 4m bgl. All of the boreholes were installed with gas monitoring standpipes.

Six rounds of gas monitoring visits were undertaken between 13 December 2011 and 2 March 2012 under varying weather conditions including falling atmospheric pressure.

The gas monitoring included the monitoring of carbon dioxide, methane and oxygen measured in each borehole along with flow rate, atmospheric pressure and water level.

The maximum concentration of methane recorded was 0.2% (LP1) and the maximum concentration of carbon dioxide recorded was 5.4% (LP8), with a maximum borehole flow rate recorded at 1.1 litres/hour (LP7, LP13 and LP14).

Through the development of a GSV in accordance with CIRIA 665 the sites highest methane concentration and the highest gas flow rate gave a GSV for methane of 0.0022. The same exercise for carbon dioxide gave a GSV of 0.0594.

Although the maximum recorded concentration of carbon dioxide marginally exceeds the TMC between "Green" and "Amber 1", it was noted that this was on one occasion only on six visits with twenty monitoring points and as the maximum flow rates recorded were typically about 1 l/hr it was concluded that a "Green" classification should be applicable to the northern part of the site and no gas protection would be required in new build in respect of methane or carbon dioxide in near surface soils.

The letter report did highlight that the site levels and soil conditions could be subject to change depending on the material movement and final levels of the proposed development.

2.9.7 17th May 2012, GEG Geo Environmental Group, Intrusive Assessment of Canal Base, Oxford Canal, Banbury, Oxfordshire

Geo Environmental Group (GEG) was commissioned to undertake an intrusive investigation of the canal base at a single location on the Oxford Canal in order to provide relevant information with respect to the proposed drilling of a 300mm drain directly beneath the canal.

The purpose of this report was to determine:

- the natural ground conditions likely to exist beneath the canal; and
- the thickness of 'puddle clay' sealing the base of the canal.

The site investigation comprised the drilling of one window sample borehole to a depth of 6.00m at the side of the canal, and the percussive hand drilling of one borehole utilising a 38mm sampler tube in the base of the canal.

The ground conditions encountered in the window sample borehole next to the canal (BH1) included a layer of re-worked topsoil which comprised red brown, slightly clayey to clayey topsoil with occasional charcoal fragments to 1.20m bgl over alluvium to the base of the borehole (6.00m).

The alluvium consisted of a layer of soft to firm, red brown, slightly friable clay (1.00m thick) overlying loose light red brown, very clayey, fine to medium sand (0.40m thick) which overlaid loose, light red, gravelly, fine to medium sand (sub-angular gravel becoming rare with depth) to the base of the borehole. SPT N-values recorded within the alluvium range from 8 at 3m bgl 14 at 6m bgl.

The Charmouth Mudstone strata were not encountered in this investigation.

During the drilling of the borehole, groundwater was encountered initially in the stratum at 2.20m (wet ground) which became very wet from 3.70m. On completion of the borehole, the groundwater was dipped at a depth 1.80m bgl.

2.9.8 August 2012, Wardell Armstrong LLP, Remediation Ground Investigation Report, Bankside, Banbury

This report was prepared to provide an assessment of the ground conditions which exist across the Bankside site and to identify potential geotechnical constraints to the proposed residential development. The works were aimed at determining the geotechnical characteristics of the ground conditions for the outline recommendation of foundation zoning areas across the site.

The investigation works comprised the excavation of 64 trial pits and the drilling of 29 light percussion probe holes and 7 cable percussion boreholes with 10 CBR tests. The investigation works were undertaken across the whole of the site area focussing particularly on the areas of the proposed residential development within the north east and south of the

site. Investigation was undertaken on an approximate 70m grid across the development areas with an approximate 200m grid across the areas of public open space.

Standpipes were installed within three of the light percussion probe holes and within two of the cable percussion boreholes to monitor groundwater levels within the shallow soils.

Relatively shallow groundwater levels were encountered and recorded in the vicinity of three springs at the site (near LPs 107, 108 and 109). Development in these areas may, therefore, require some groundwater control measures to keep excavations dry and precautions to maintain stability. In the historic site investigations groundwater was encountered within three of the cable percussion boreholes, five of the trial pit excavations and 11 of the light percussion probe holes across the site area. As a generality the groundwater was encountered in four main zones as follows.

- Groundwater was encountered within the south west of the site adjacent to Oxford Road where the investigation positions (LP123, LP125, TP162 and BH105) penetrated the Marlstone Rock Bed strata into the Dyrham Formation. The water was noted to be a slight seepage at approximately 2.0m depth with a greater flow noted at a depth of approximately 5.0m below ground level.
- Groundwater was noted within the west of the site within the field adjacent to Canal Lane within investigation positions LP115 and LP116 as damp soils materials at approximately 1.5m depth below ground level. This is conjectured to be within the Whitby Mudstone Formation and was noted to be directly above the weathered sandstone materials.
- Investigations were undertaken within the vicinity of three springs recorded on the historical mapping of the site. LP107, LP108 and LP109 all recorded groundwater seepages at depths of between 1.03m and 2.0m depth below ground level. This corresponds with the conjectured spring locations.
- Groundwater was also noted within the north east of the site close to the Oxford Canal. Two light percussion probe holes, two cable percussion boreholes and four trial pits within this area record ingress of groundwater during the undertaking of the investigation. It is conjectured that the groundwater levels encountered within these positions represents the water table within this area. The groundwater was encountered at depths of between 0.9m and 2.9m within these materials although it is considered that the groundwater encountered within TP107 at 0.9m depth may represent a perched water level within sand lenses present within the clay materials. It is therefore considered that the water table within this area of the site is approximately 1.5m to 2.5m depth below the surface level.

Selected samples of soils from the trial pits and cable percussion boreholes were analysed to determine their geotechnical characteristics.

The laboratory testing comprised the following:

- 37 samples for natural moisture content;
- 25 samples for plastic limit and liquid limits;
- 34 samples for particle size distribution;

- 20 samples for pH and sulfate content; and
- 4 samples for compaction with a 4.5kg rammer.

No undisturbed samples of material were retrieved as part of the site investigation due to the nature of the ground conditions at the site. It was, therefore, not possible to undertake compressibility or shear strength testing on undisturbed samples of materials.

The average moisture content within the cohesive materials was found to be 24% with the average moisture content within the granular materials of 23%.

The modified plasticity index results range from 11% to 49% and were split into the following groups pertaining to the underlying geology at the site:

- Charmouth Mudstone Formation, moderate to high volume change potential;
- Marlstone Rock Bed, low volume change potential; and
- Whitby Mudstone Formation, medium volume change potential.

Similarly the particle size distribution results were split into the following groups pertaining to the underlying geology at the site.

- The Charmouth Mudstone Formation is silty and sandy clay materials with some gravelly clay materials. The fines content ranges from 39% to 96% although the majority comprise over 51% fines.
- Only one sample of material from the Dyrham Formation was tested, with a fines content of 35% and is described as gravelly, sandy clay.
- The Marlstone Rock Bed is clayey, sandy gravel and cobbles to very gravelly clay. The analysis records the gravel and cobble content between 20% and 84% although the majority, are greater than 30%.
- The Whitby Mudstone Formation is dominated by clay and silt which is likely to be present as a thin band above the Marlstone Rock Bed in the area.

Four samples were used to determine the dry density / moisture content relationship. Two were from the proposed development area within the north east of the site and two from the south west of the site. The results indicate that the materials are generally wet of the optimum moisture content (OMC) and the maximum dry densities vary between 1.63Mg/m³ and 1.90Mg/m³ with OMC between 14% and 21%.

The CBR results indicate that whilst the shallow ground within the south of the site recorded CBR values of between3% to 12%, the shallow ground conditions close to the Oxford Canal indicated softer materials with CBR values ranging from 0.9% to 1.6%.

It was highlighted that detailed pavement design should take account of these results and appropriate measures be taken in the construction of the roads.

The geotechnical assessment was undertaken based on the information from both the 2012 site investigation works and the previous investigation phases. The assessment identified three generalised foundation zones.

These three areas comprise:

- Zone A (the area of residential development in the northern part of the site with in the former tip area) – variable foundation solutions including possible vibro-stone or piled foundations within the clay and sand-based Made Ground materials. Foundation loads would either need to be transferred to the underlying stiff clay materials or the bearing properties of the Made Ground materials improved;
- Zone B (the area of residential development in the northern part of the site that is outside the tip area) shallow strip foundations within the firm to stiff, grey clay materials at depths of approximately 1.0m depth below ground level. An allowance should be made for possible localised soft materials where soft materials may require removal and replacement with more competent materials or concrete / trench fill; and
- Zone C (the area of residential development in the south part of the site) shallow strip foundations may be constructed within the stiff clay materials or within the weathered sandstone at shallow depths across this section of the site. An allowance should be made for treatment of localised soft spots if encountered. Investigation undertaken within this zone has also indicated localised soft materials to a depth of 2.7m bgl therefore foundations loads would need to be transferred to the underlying stiff clay materials.

2.10 Suitability of Previous Data

2.10.1 Geotechnical Test Data

The geotechnical testing undertaken at the site is reasonably extensive and considered suitable. However, additional trial pitting will be needed to confirm the findings for design purposes. The information will be incorporated into the assessment of the site undertaken within this report and will enable an appropriate zoning of foundation solutions.

2.10.2 Chemical Test Data (Soil)

The chemical testing undertaken at the site is limited in its spatial distribution and selection of determinands. The chemical test data are, however, useful and will be incorporated into the data set used in the assessment of the site undertaken within this report.

2.10.3 Chemical Test Data (Water)

Only limited groundwater/surface water has been encountered and sampled at the site to date. The testing of water and leachate samples from the site undertaken to date does not indicate a significant potential for contamination.

2.10.4 Ground Gas Data

There are substantial amounts of ground gas data available for the north part of the site which is in accordance with current guidelines (CIRIA 665). Hydrock considers the data to be suitable for the purposes of this report.

3.0 ADDITIONAL GROUND INVESTIGATION

3.1 Investigation Rationale

The ground investigation rationale was to confirm the findings of previous work and fill in gaps of the existing ground investigation information at the site as summarised in Table 3.1.

Table 3.1: Investigation Rationale

Exploratory Holes	Purpose		
Tip Area in the north of the site			
HTP 01-06	To investigate the depth and physical and chemical composition with a view to the foundations requirements and to provide further samples for chemical testing.		
Remainder of site			
HTP 11-46	To assess shallow ground conditions with a view to the foundation requirements and to provide further samples for chemical testing.		

3.2 Site Works

The position of site investigation locations (surveyed using GPS) are shown on the Ground Investigation Plan (Drawing C12702-G002) in Appendix C.

The site works undertaken for this investigation are summarised in Table 3.2. The logs are presented in Appendix C.

Table 3.2: Summary of Site Works

Activity	Method	No.	Max. Depth (m)
Trial pits	Machine (JCB 3X)	46	3.5

3.3 <u>Geo-Environmental Testing</u>

3.3.1 Sampling Strategy and Protocols

The locations of the investigatory holes were located to provide a reasonable spread of information on the site ground conditions and to fill in the gaps in the coverage of the previous site investigation information. No specific sampling statistics or grid were utilised in this instance.

Samples were taken stored and transported in general accordance with BS 10175:2011.

3.3.2 Laboratory Analyses

The findings of the former site investigation and risk assessment have been used to scope the analyses of chemicals of potential concern as follows.

The following were performed on samples of **soil or other solids**:

 25 Hydrock default suite of determinands for solids comprising: As, B (water soluble), Be, Cd, Cr (III), Cr(VI), Cu, Hg (inorganic), Ni, Pb, Se, V, Zn, cyanide (free), pH, asbestos screen, speciated polycyclic aromatic hydrocarbons (PAH, by GC-MS), total phenols and fraction of organic carbon; and 4 Hydrock default waters suite of determinands, following leaching to BS12457-2, comprising: Ag, Al, As, B, Ba, Cd, Co, Cr (III), Cr(VI), Cu, Fe, Hg, Mn, Mo, Na, Ni, Pb, Sb, Se, Sn, Zn, V, cyanide (total), phenols (total), ammonium, bromate, chloride, fluoride, nitrate, nitrite, sulfate, PAH (speciated), pH, EC and hardness.

The chemical test results are provided in Appendix E.

4.0 GROUND INVESTIGATION DATA

4.1 Physical Ground Conditions

The ground conditions encountered during the current works are in general accordance with the expectations from the previous investigation works at the site.

In 2012 Hydrock were not given permission to access a paddock and large field at the southern end of the site and as a result the proposed trial pits in this area had to be abandoned. These are HTP30, HTP31, HTP34, HTP36 and HTP37 which accounts for their absence from this report.

In 2013, when access was permitted Hydrock progressed a total of six trial pits within this area, these are labelled HTP41 to HTP46.

Details are provided in the logs in Appendix C and the individual strata are described in the sections below. The geology of the site is shown on the Geo-environmental Zonation Plan (Drawing C12702-G003) in Appendix A.

4.1.1 Topsoil

Topsoil was encountered across the site. It is a sandy, silty, gravelly clay or gravel in the southern area underlain by the Whitby Mudstone Formation and Marlstone Rock Bed, and a sandy, silty, gravelly clay in the central and eastern areas underlain by the Dyrham and Charmouth Mudstone Formations. The tip in the north has a partial covering of brown, clayey, sandy topsoil.

4.1.2 Made Ground

Made Ground was only encountered in the tip area in the north part of the site, at depths ranging from the ground surface to 1.6m bgl.

In general, the Made Ground is of variable content comprising gravelly clay with cobbles and boulders, gravel and cobbles and boulders of brick, white polystyrene, plastic sheeting, concrete, concrete fragments, plastic. A boulder-sized section brick walling was encountered.

4.1.3 Charmouth Mudstone Formation

The Charmouth Mudstone Formation was encountered in the northern half of the site including underlying the Made Ground, at depths ranging from 0.2 to 1.6m bgl. The Charmouth Mudstone Formation consisted of low to high strength, orange brown, brown and grey, silty residual clay and moderately packed, light orange brown, residual, fine sand and with depth firm dark brown, mottled, orange brown and grey, residual, clay with frequent fine to medium extremely weak lithorelicts (encountered from 1.7 to 2.2m bgl).

At the lower end of the slope in the northern part of the site the Charmouth Mudstone encountered in HTP04 and HTP06, adjacent to the Oxford Canal, was unstable, moderately packed, brown and orange brown, fine sand and moderately packed, red brown, clayey, gravelly, fine sand with groundwater seepage (encountered between 0.9 and 1.8m bgl).

4.1.4 Dyrham Formation

The Dyrham Formation was encountered in HTP12 in the centre of the site and consisted of very low to low strength, orange brown and grey, silty, residual, wet clay to a depth of 2.8m bgl where the trial pit was terminated because of collapse.

4.1.5 Marlstone Rock Bed

The Marlstone Rock Bed was encountered in the southern half of the site. This generally consisted of medium to high strength, red brown and brown, sandy, gravelly, residual clay over moderately strong, blue grey and red brown banded, shelly, partially weathered limestone.

4.1.6 Whitby Mudstone Formation

The Whitby Mudstone Formation was encountered in the southern end of the site and consisted of a layer of medium to high strength, firm to stiff, brown and yellow brown mottled, gravelly, sandy, silty residual clay which was underlain by weathered limestone of the Marlstone Rock Bed.

4.2 <u>Geo-Environmental Results</u>

Soil Chemistry

The chemical test results are given in Appendix E.

Concentrations of metals and organic compounds (PAHs) were detected above the laboratory detection limit in all soil samples.

As indicated in previous investigations, notable concentrations of arsenic were recorded across the site and notable concentrations of organic contaminants encountered in the northern part of the site associated with the Made Ground. Vanadium was also detected in significant concentrations, although this was not tested for by previous investigators.

It is apparent from the arsenic and vanadium distribution that there are two distinct geochemical provinces related to the underlying geology: the area underlain by the Marlstone Rock Bed and Whitby Mudstone Formation having higher concentrations than the area underlain by the Dyrham and Charmouth Mudstone Formations.

Asbestos

The presence of asbestos was not detected during laboratory analysis on the soil samples. However, due to the uncontrolled nature of the tip the presence of asbestos cannot be discounted.

Leachate Chemistry

Eluate (leachate) testing was undertaken to assess the risk to controlled waters from soil contaminants in line with good practice defined in ISO 15175:2004.

The test results are given in Appendix E.

There is no particular spatial distribution of chemicals of potential concern that requires explanation.

5.0 GENERIC QUANTITATIVE RISK ASSESSMENTS

5.1 Conceptual Site Model

The additional site investigation has not produced any significant changes to the conceptual site model which is summarised as follows.

The site has not been previously developed and has a history of agricultural use. However, it is known that the northern site area has previously been used as a tip. Therefore the site can be split between the tip area and the wider site area.

For the wider site area, testing of samples of natural materials from the site indicates that there are two geochemical provinces: the Marlstone Rock Bed and Whitby Mudstone Formation having higher metal concentrations than the Dyrham and Charmouth Mudstone Formations.

The tip area has been shown to include various types of Made Ground. From the investigation and testing these materials have been found to contain elevated contaminants recorded in various locations throughout the tip.

Only limited groundwater/surface water has been encountered and sampled at the site to date. The south end of the site which overlays the Marlstone Rock Bed is a Secondary A Aquifer and springs are observed at the eastern end of the site which are likely to expel where the Marlstone Rock Bed changes to the either the Dyrham Formation or the Charmouth Mudstone Formation.

The tip area is not underlain by the Secondary A Aquifer and is considered unproductive strata.

The Oxford Canal forms the north eastern site boundary and the River Cherwell is located approximately 230m further to the north.

The plausible contamination linkages are:

- risks to the health of end users of the site from substances in the natural soils and Made Ground;
- risks to plant life from metals in the natural soil and Made Ground; and
- pollution of the surface water environment by runoff of contaminants in the Made Ground.

These linkages are assessed in the following sections.

Generic risk assessment is a two stage process. Firstly, the measured contaminant concentrations are compared to the relevant GACs. This is the Risk Estimation stage. Where there is a suitable dataset, this is done after carrying out statistical analysis to determine the upper confidence limit on the true mean. Otherwise, maximum or specific data points are compared directly.

The second stage, Risk Evaluation, comprises an authoritative review of the findings with other pertinent information, in cases where the GACs are exceeded, in order to consider if exceedance may be acceptable in the particular circumstances.

5.2 Human Health and Plant Life

This is a Tier 2 assessment using soil screening values and involves generic human health risk assessment for the CLEA **residential with plant uptake** land use scenario.

The soil chemical analysis results from this ground investigation and historical ground investigation information have been screened against guideline soil concentrations to provide an assessment of potential risks associated with contamination at the site. Justification for the criteria adopted for this risk assessment is given in Appendix D.

It has been assumed in this report that the exposure conditions are within the generic conditions used to derive Soil Guideline Values (SGVs). Where no SGVs are published, or where the published values require modification, a number of Generic Assessment Criteria (GAC) have been developed for certain chemicals of potential concern. These GAC have been developed using largely generic assumptions about the characteristics and behaviour of sources, pathways and receptors, i.e. similar to those used by the Environment Agency in the derivation of SGVs.

It should be noted that the term "further assessment required" is used to denote soil concentrations that are equal to, or exceed, a GAC. This does not automatically mean that the soil is "contaminated".

5.2.1 Risk Estimation (Including Statistical Testing)

The 'averaging areas' used in this report are based on the conceptual model and the proposed development and are summarised as:

- 1. the former tip area in the north part of the site which is proposed for residential development; and
- 2. the wider site area underlain by natural ground of the Marlstone Rock Bed and Whitby Mudstone Formation.

Initial Data Review

The data set for each chemical determinand has been assessed for the presence of potential outliers (based on the conceptual model) and to determine if the data are normally or non-normally distributed, in line with the methods described in Appendix D.

No outliers have been removed.

Statistical Testing

Suitable data sets are available for statistical analysis of the determinands tested.

Using the methodology detailed in Appendix D and in line with the guidance provided by the CIEH (May 2008) the 95^{th} upper confidence level on the true mean (US₉₅) has been calculated from the sample data.

Data have been assessed using the one-sample t-test (for data which can be treated as normally distributed or that do not deviate significantly from normal) or the one-sided Chebychev Theorem (for significantly non-normally distributed data).

Appendix E contains the detailed results of the statistical assessment for each chemical of potential concern, together with summary sheets for human health and plant life. The results are summarised below.

Human Health & Plant Life

With regards to human health, based on a US_{95} exceedance of the GAC, the pervasive chemicals of potential concern which require further assessment and/or remediation are summarised in Table 5.1 and Table 5.2 below.

Table 5.1: Pervasive Chemicals of Potential Concern for Which Further Assessment is
Required (Human Health)

Chemical of Potential Concern	Generic Criterion (mg/kg)	Basis for Generic Criterion	No. Samples	Min. (mg/kg)	Max. (mg/kg)	No. Samples Exceeding Generic Criterion	US ₉₅ (mg/kg)
Tip Area		•					
Arsenic	32	SGV report + CLEA 1.06	28	12	150	19	80
Vanadium	74	LQM/CIEH + CLEA 1.06	7	0.5	230	4	230
Benz(a)anthrace ne	4.7	LQM/CIEH + CLEA 1.06	19	0.031	36	4	16
Benzo(a)pyrene	0.94	LQM/CIEH + CLEA 1.06	19	0.017	56	9	20
Benzo(b)fluorant hene	6.5	LQM/CIEH + CLEA 1.06	19	0.047	40	4	16.2
Chrysene	8	LQM/CIEH + CLEA 1.06	19	0.027	40	4	18
Dibenz(a,h)anthr acene	0.86	LQM/CIEH + CLEA 1.06	19	0.01	4.5	4	1.8
Indeno(1,2,3,cd) pyrene	3.9	LQM/CIEH + CLEA 1.06	19	0.01	43	4	15
Naphthalene	3.7	LQM/CIEH + CLEA 1.06	19	0.01	42	3	14
Wider Site Area Underlain by Marlstone Rock Bed and Whitby Mudstone Formation							
Arsenic	32	SGV report + CLEA 1.06	40	17	230	37	111
Vanadium	74	LQM/CIEH + CLEA 1.06	16	100	740	16	550

Table 5.2: Pervasive Chemicals of Potential Concern for Which Further Assessment is Required (Plant Life)

Chemical of Potential Concern	Generic Criterion (mg/kg)	Basis for Generic Criterion	No. Samples	Min. (mg/kg)	Max. (mg/kg)	No. Samples Exceeding Generic Criterion	US ₉₅ (mg/kg)
Wider Site Area Underlain by Marlstone Rock Bed and Whitby Mudstone Formation							
Nickel	75	BS3882 2007	40	29	210	31	127
Zinc	300	BS3882 2007	23	74	970	5	394

5.2.2 Risk Estimation (Without Statistical Testing)

In this section the results are discussed for determinands for which the data set does not allow application of statistical analysis by virtue of low sample numbers. This applies to the wider site area underlain by natural ground of the Dyrham and Charmouth Mudstone Formations.

Reference to the summary sheets in Appendix E shows that there are no exceedances of the GACs for human health or plant life.

5.2.3 Risk Evaluation – Human Health

Tip Area – Made Ground

The metals in the Made Ground are likely to have been derived from soils in the region because the tipped material appears to be construction waste incorporating a significant proportion of soil. The comments made below concerning natural soils elsewhere apply. However the risk driver is seen to be the PAHs in the Made Ground. As an example, benzo(a)pyrene is present at a concentration of 60 times the GAC and consequently the soils in this area are considered unsuitable for use as near surface soils in residential developed areas of the site and mitigation measures will be required.

Wider Site Area Underlain by Marlstone Rock Bed and Whitby Mudstone Formation

Chemical testing has indicated elevated concentrations of arsenic and vanadium above the GACs. DEFRA (2005) has made it clear that exceedance of a GAC does not necessarily meet the legal test of contaminated land, i.e. exceedance of a GAC does not necessarily equate to unacceptable risk. Consequently, the GACs must be considered as screening values only and the exceedances have been further assessed below.

Arsenic

Chemical testing has indicated pervasive concentrations of arsenic ranging from 17 to 230mg/kg with a US₉₅ percentile of 111 mg/kg within the natural soil.

There is also a naturally high background level of arsenic in the area, which ranges from 30 to 820mg/kg and, as with vanadium, this is not considered a significant risk.

In addition the six PBET test results at the site means it is possible to assess the percentage of arsenic that is bioaccessible and integrate this information into the CLEA model to derive a Site Specific Assessment Criteria (SSAC) for this area of the site.

Using the highest percentage of arsenic bioaccessibility encountered in the areas underlain by the Marlstone Rock Bed formation a worst case of 5.9% bioaccessibility has been used.

The CLEA model has therefore been used to produce SSAC. All other parameters were set to default values in the CLEA v1.06 model, run for residential with plant uptake, which generated an SSAC of 136 mg/kg. This is greater than the US_{95} which indicates no significant risk.

It is concluded that no further assessment or mitigation with regard to arsenic is required. This has been confirmed by Mr S Gregory, Environmental Protection Officer at Cherwell DC (email to Hydrock dated 28 February 2013).

Vanadium

The GAC for vanadium (74mg/kg) is exceeded in the area of the site underlain by the Marlstone Rock Bed and Whitby Mudstone Formation, with the US₉₅ being 550mg/kg.

These strata are known to have naturally high levels of vanadium, which in this area range between 122mg/kg and 626mg/kg according to the *Advanced Geochemical Atlas of England and Wales* (Rawlins *et al* 2012) which states that elevated concentrations in the UK occur mainly over Jurassic strata especially between Banbury and Melton Mobray (spatially associated with the ironstones).

According to Breward (2007) the vanadium may be a resistate element, one whose host minerals are highly resistant to both chemical and physical weathering and hence have a very long residence time in soils and sediments. However, there is a strong correlation between iron and vanadium concentration and Breward attributes this, at least in part, to *in situ* sorption and concentration by sedimentary iron oxides during their formation. It is possible that an Fe(III) vanadate complex replacement mineral related to the strengite-scorodite group is formed. The arsenic and other trace elements in the soil are similarly attributed to co-precipitation enhancement during mineral formation.

Comparing the sediment and the stream water vanadium contents, Breward concludes that because of the stability of the iron oxides, the environmental mobility and bioaccessibility of the vanadium (and arsenic) is extremely low.

Whichever of these mechanisms is the actual one operating, this means that the vanadium is tightly bound in the mineral crystal lattice and has survived as such for millions of years. These strata are of Toarcian age, which spans the time between 183.0 and 175.6 million years ago.

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The GAC report (Nathanail *et al* 2009) states that naturally occurring vanadium in soils is generally relatively insoluble V(III), but can be present in the more soluble V(V) form. The bulk of and released vanadium is retained in the soil, associated with the organic matter because humic acids convert the more mobile meta-vanadate anions into the immobile vanadyl cations resulting in local accumulation. There may be some mobility under oxidising, unsaturated conditions, but the vanadium is immobile under reducing, saturated conditions.

Consequently, this exceedance is not likely to constitute a significant risk to human health, in line with the current Contaminated Land Statutory Guidance, which accepts that there may be natural background levels of substances as a result of geology.

Further consideration of the likely risks can be undertaken by way of a sensitivity analysis of the CLEA 1.06 derived GAC. The GAC is an initial screening value and not a remediation standard and as such is very conservative. It is designed to protect against all forms of vanadium, most particularly the most common commercial form, vanadium pentoxide, which is a strong oxidising agent and has many important industrial uses.

The GAC report (Nathanail *et al* 2009) states that vanadium occurs in six oxidation states and has a complex chemistry, forming more compounds than any element except carbon. The toxicologically most significant ones are V_2O_5 , NaVO₃, Na₃VO₄, VOSO₄ and NH₄VO₃. The toxicity increases with the oxidation state (Expert Group on Vitamins and Minerals 2003) and the toxic effects are mostly the result of inhalation, with very few reported cases relating to ingestion. In fact, vanadium is used in dietary supplements as VOSO₄ and NaVO₃ up to 0.025mg/day, although there is no evidence to support any useful effect. According to the Oak Ridge National Laboratory (Opresko 1991) V(V) is five times as toxic as V(III).

The Health Criterion Values (HCV) used in the CLEA model are designed to protect against the simple, and much more toxic, vanadium compounds. The complex vanadium-iron minerals of the soils beneath the site will display significantly reduced toxicity.

There is no information available, however, that can be used to redefine the HCV with respect to these minerals and so the default values are retained in this exercise.

An important parameter in the derivation of the GAC is the soil-water partition coefficient (Kd). This dictates how much of the vanadium stays in the soil and how much can go into solution, thereby making it mobile and capable of entering the vascular system in the body. The range of values reported in the GAC report, Section 7.7.3, is 12.6-1000 l/kg. One of the studies cited was by the USEPA (2005) which reported a range of 3.2-316, with a mean of 50.1 l/kg. The lowest value was adopted in the derivation of the GAC as the default because of this high range the authors wished to be as conservative as possible.

There is insufficient data available to derive a site-specific Kd value, so a range of values has been tried in the sensitivity analysis.

The highest solubility in water was adopted by the GAC authors (Section 7.7.2 of the report) as a conservative value. However, sensitivity analysis has shown the GAC to be insensitive to solubility in this instance and the default value has been retained.

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As mentioned by Breward above, the bioaccessibility is much lower than the 100% assumed as a default in the CLEA model. This is mentioned by a number of sources, including Oak Ridge National Laboratory which reports 0.5-2% for ingestion and 20-25% for inhalation. The maximum values have been adopted in this exercise.

Accordingly, the CLEA model has been run setting the oral and inhalation relative bioavailabilities to 0.02 and 0.25, respectively and varying the Kd. In order to use the Kd in the model, the soil to plant concentration factors are changed from the default numerical values in the GAC report to those calculated internally by the CLEA model. This has been verified by running the model with all the vanadium defaults and only changing these factors, and there is no change to the residential GAC.

The results of the sensitivity analysis for Kd are as follows:

	Residential SSAC (mg/kg)
Kd (l/kg)	(Two significant figures)
12.6 (default)	120
45	380
48	410
50	420
60	490
69	550
70	560
100	730

It can be seen that for the SSAC to be the same as the US_{95} , the Kd need be only 69 l/kg and this is well within the range of literature values of 12.6-1000 l/kg and is close to the mean value quoted by the USEPA.

Given the published comments concerning the stability of the iron-vanadium minerals, it is entirely feasible that the Kd will be 69 l/kg or greater.

Furthermore, the US_{95} of 550mg/kg is based on the use of the non-parametric statistical test (Chebychev Theorem) as a conservative measure. The visual assessment of the normality of the data set distribution is included in Appendix E as a Q-q plot and histogram. Although the data are non-normal, the degree of non-normality is not great and it is highly likely that the parametric statistic (t-test) could cope (Barnes *et al* 2010). A check using the Shipiro-Wilk test for normality shows the data set to be just outside the definition of normally distributed because the W statistic is 0.813, a only little below the W-critical value of 8.887.

In which case, the US $_{95}$ would be 405mg/kg and the required Kd would be 48 l/kg, less than the USEPA (2005) mean value.

Given that such a large swathe of the country is underlain by sediments of this nature, there will be a great number of developments where no mitigation measures have been incorporated. Indeed, it is believed that the current site use includes the commercial

production of vegetables. Hydrock is not aware of any evidence to suggest that existing residents are being affected by the presence of naturally occurring vanadium.

It is concluded that the presence of naturally-occurring vanadium is not a significant risk to human health.

This assessment has been presented to Cherwell DC for consideration, but no reply has been received at the time of writing.

5.2.4 Risk Evaluation – Plant Life

Plant Life

With regards to plant life the testing and statistical assessment to date indicates apparently pervasive nickel and zinc with US_{95} of 127mg/kg and 394mg/kg versus GAC of 75mg/kg and 300mg/kg, respectively. With reference to *Advanced Geochemical Atlas of England and Wales* (Rawlins *et al* 2012), high zinc and nickel are expected within the Marlstone Rock Bed.

Whilst detriment to plant life is hard to quantify as many of the GACs are based on agricultural crop yields rather than serious harm of death of a species. The exceedance for zinc is slight and that for nickel is not large. Given that the vegetation on site did not show any signs of physical distress and the land is currently use for farming, Hydrock does not believe any additional consideration is required with regards to risks to plant life.

5.3 Pollution of Controlled Waters

5.3.1 Risk Estimation

The risks to groundwater and surface water have been assessed according to the remedial targets methodology (RTM) prescribed by the Environment Agency (2006) as described in Appendix D. Pollutant inputs from contaminated land sites are considered as passive inputs under the European Water Framework Directive (2000/60/EC) (WFD) and its daughter Directives, and as such are regulated under the Agency's 'limit' pollution objective.

Acceptable water quality targets (WQT) are defined for protection of human health (based on drinking water standards (DWS)) and for protection of aquatic ecosystems (environmental quality standards (EQS)). For the purposes of this report, the site data are compared with the various targets as set out according to the Hydrock scenario(s) in Table 5.2 (see Appendix D for details), on the basis of the following:

- the tip area in the north of the site is underlain by the Charmouth Mudstone Formation which is classified as unproductive strata and the Oxford Canal is adjacent to the east of the tip area, therefore there is a low risk of leachate entering the Oxford Canal through surface water runoff or leachate migration; and
- the southern part of the site underlain by the Marlstone Rock Bed and Whitby Mudstone Formation is underlain by a Secondary A Aquifer and therefore there is a low risk of leachate entering the aquifer and migrating to the Oxford Canal to the east of the site.

Hydrock Scenario	Water Body Receptors	Secondary Receptors	Example Contaminant linkages	RTM Level and Data Used	Water Quality Targets
Tip area	<u>.</u>				
F	Surface water.	Aquatic ecosystem.	Contaminants from site leach or seep into surface water which may be an aquatic ecosystem.	RTM Level 1 - Soil leachate.	EQS (inland)
Wider Site A	rea				
В	Groundwater. Surface water.	Aquatic ecosystem.	Contaminants from site leach or seep into groundwater body and this feeds surface water by base flow. The surface water may be an aquatic ecosystem.	RTM Level 1 - Soil leachate.	EQS (inland)
pathway), or i	s estimated from	national maps.	his is measured either in the receiving wate	C (it is part of the

Inland waters EQS applicable to freshwater, other waters EQS applicable to marine or transitional waters.

Where both DWS and EQS are applicable, it is assumed that the EQS is for inland waters.

This table and the results of the assessment are considered as a first screening for potential risks of pollution of Controlled Waters. More specific requirements may be stipulated by the Environment Agency.

The results of the remedial targets methodology assessment are presented in Appendix E and are summarised in Table 5.3.

It should be noted that in some instances the reporting limit (or detection limit) quoted by the laboratory may be greater than the water quality target that it is being assessed against. As the current exercise is an initial screening assessment, further assessment of these elements has not been undertaken if all the results for all samples are below the detection limit but above the water quality target. However, in other cases, even though the detection limit is greater than the water quality target, some sample results do exceed the target and assessment is viable. See Appendix D for comment on detection limits.

Chemical of Potential Concern	Water Quality Target (ug/l)	Basis for Water Quality Target	No. Samples	Min. (ug/l)	Max. (ug/l)	No. Samples Exceeding Target
Tip Area						
Cu (dissolved)	6	Inland Water EQS	2	1	6.3	1
Benzo(a)pyrene	0.05	Inland Water EQS	2	6.7	8.9	2
PAH sum of benzo(b)fluoranthene benzo(k)fluoranthene	0.03	Inland Water EQS	2	0.02	1.21	2
Fluoranthene	0.1	Inland Water EQS	2	0.01	0.6	1
Naphthalene	2.4	Inland Water EQS	2	2.6	3.3	2
Wider Site Area						
Fe (dissolved)	1000	Inland Water EQS	20	1	1200	1
Zn (total)	8	Inland Water EQS	21	1	27	4
Benzo(a)pyrene	0.05	Inland Water EQS	12	0.02	7.1	2

Table 5.3: Chemicals of Potential Concern for Which Further Assessment is Required
(Controlled Waters)

Chemical of Potential Concern	Water Quality Target (ug/l)	Basis for Water Quality Target	No. Samples	Min. (ug/l)	Max. (ug/l)	No. Samples Exceeding Target	
Tip Area							
Fluoranthene	0.1	Inland Water EQS	12	0.01	1	1	
Cr(iii)(diss)	417	Inland Water EQS	15	1	8	2	

Note: the 95% ile value is compared with the water quality target and a June 2007 Site Investigation of the Spring and Surface Water demonstrated no exceedances of the Inland Water EQS.

* The Water Supply Regulations 1989 and the Private Water Supply Regulations 1991 both contained a prescribed concentration of 10 µg/l for "dissolved or emulsified hydrocarbons (after extraction with petroleum ether); mineral oils". This was removed when these Regulations were updated in 2000 (consolidated 2007) and 2009, respectively. However 10 µg/l is used in this report as an initial screening assessment as it is frequently the preferred approach of the Environment Agency.

5.3.2 Risk Evaluation

Tip Area – Made Ground

The data indicate that the EQS are exceeded for copper and various PAH species.

The surface gradient is to the eastwards to the Oxford Canal but no groundwater was encountered within the Made Ground with the exception of a slight seepage of perched water in HTP3.

In general, it can be concluded that the Made Ground within the tip area is contaminated with leachable copper and PAHs. However it is that unlikely that contaminants are significantly mobile due to the high clay content of the Made Ground and underlying solid geology and any lateral movement away from the Made Ground would degrade due to natural attenuation of the of the leachate. In addition the proposed development will add hard standing and soil cover that will further limit the potential for the infiltration of water into the Made Ground and further minimise the potential migration of contaminants within the Made Ground.

Whilst there are elevated concentrations of Chemicals of Potential Concern, based on the investigation works undertaken to date and subject to agreement with the Environment Agency, Hydrock does not believe the site poses a significant risk to Controlled Water for the following reasons:

- the high clay content of the Made Ground and underlying Charmouth Mudstone Formation forms an aquiclude to prevent migration of leachate vertically and laterally to the Oxford Canal, which is not at risk;
- the shallow waters at, and in the vicinity of, the site are not abstracted for human consumption;
- whilst the ecosystem in the Oxford Canal to the east of the tip area is a potential receptor, the reduction of concentrations due to leachate attenuation will reduce the risks as the distance from the Made Ground increases. In addition the Wardell Armstrong 2007 site investigation tested a sample of water associated with the canal and no exceedances of the GAC were noted; and
- subject to a piling risk assessment showing no additional pathways to the Oxford Canal, no changes to the groundwater regime are expected to occur during development.

Wider Site Area – Natural Ground

The data indicate that the EQS are exceeded for metals (chromium (III), iron and zinc) and PAH species (benzo(a)pyrene and fluoranthene) leachate within the natural ground.

The general groundwater gradient is to the east and follows the surface contours. The groundwater in Marlstone Rock Bed provides base flow to the springs in the east of the site. A limited suite of metals were determined in the 2007 investigation at the site, in two spring water samples. The current (2013) EQS for copper was exceeded in one of these samples.

There is no obvious source of PAHs in the natural soil and the most likely explanation is fallout from atmospheric pollution.

Whilst there are elevated concentrations of Chemicals of Potential Concern, based on the investigation works undertaken to date and subject to agreement with the Environment Agency, Hydrock does not believe the site poses a significant risk to Controlled Water for the following reasons:

- the elevated metals are likely attributed to the underlying geology of the Marlstone Rock Bed and are typical of the wider area; and
- the PAHs are likely to be derived from atmospheric pollution and not a source of PAHs within the soil.

5.3.3 Radon

Reference to the Annex A maps in BR 211 (Scivyer 2007), based on the Indicative Atlas of Radon in England and Wales (Miles *et a*l 2007) indicates that **full** radon protection is required for new dwellings at this location in line with current guidance.

5.4 Summary of Findings of the Generic Risk Assessments

Particular areas of the site which are of potential concern are indicated on the Geoenvironmental Zonation Plan (Drawing C12702-G003) in Appendix A.

Receptor Group	Unacceptable Pervasive Pollution Source
Human Health	PAHs and potential asbestos containing materials in the Made Ground associated with the tip area in the north of the site. Elevated naturally occurring vanadium in the areas underlain by the Marlstone Rock Bed and Whitby Mudstone Formation are not considered to represent a risk to human health, but this has yet to be confirmed by the Local Authority.
Human health / property (from ground gases)	Full radon protection is required for new dwellings at this location in line with current guidance.
Controlled Waters	None
Plant Life	None

Table 5.4: Summary of Unacceptable Contaminant linkages

6.0 GEO-ENVIRONMENTAL CONCLUSIONS AND RECOMMENDATIONS

6.1 <u>Human Health</u>

Approximately 3.3ha of the northern site area is known to have been used to deposit Made Ground where trial pits have recorded up to 4m of builders' waste in a gravelly, sandy, clayey matrix.

The risk assessment undertaken in Section 5.2 indicates that the Made Ground constitutes an unacceptable risk from PAHs. Asbestos containing materials were encountered in a previous investigation. This area will require remediation / mitigation prior to redevelopment for residential end use.

Proposed mitigation options include:

- removal of Made Ground, and / or
- installation of a clean cover over remaining areas of Made Ground.

One on the contaminants of concern is naphthalene, which is semi-volatile and smells of moth balls. It is not possible to say if a cover on its own would severe the indoor vapour pathway and numerical modelling is notoriously uncertain. Even if there are no adverse health effects, there may be residual nuisance odours. For these reasons the addition of a vapour membrane in buildings is recommended. However, since full radon protection measures are also required, it will be possible to design the radon barrier to also prevent the ingress of naphthalene vapours.

Outline mitigation measures are discussed in Section 6.6.

6.2 <u>Controlled Waters</u>

The Level 1 risk assessment undertaken in Section 5.3 indicates that whilst there are elevated concentrations of Chemicals of Potential Concern Hydrock does not believe the site poses a significant risk to Controlled Waters. Consultation with regulators is recommended to confirm this.

6.3 <u>Construction Materials</u>

6.3.1 Water Pipelines

The current guidance on selection of materials for water supply pipes to be laid in contaminated land is contained in UKWIR Report 10/WM/03/02 (re-issued 2010) which sets out in Table 3.1 of that document threshold values for a selection of organic contaminants that may have a detrimental effect on pipes and fittings. However, the document is for guidance and is not mandatory and has not been adopted universally by all water suppliers.

In addition, various consultative technical bodies have expressed concern on the nature of the document and the methodologies proposed, which would result in significant cost and time implications for all site assessments. It is Hydrock's opinion that the guidance is not appropriate and it has not been followed as part of this report.

However, the findings of this investigation have been compared to the threshold values in UKWIR Table 3.1 as far as is practicable to give an indication of the possible restrictions to the use of plastic water pipes.

The majority of the site is greenfield and the investigation and assessment has indicated elevated concentrations of naturally occurring arsenic, vanadium associate with the underlying Marlstone Rock Bed. It is envisaged that standard pipework will be suitable for the site. However, this investigation was not designed specifically for water pipe runs and because of conflicting and ambiguous guidance, confirmation should be sought from the water supply company at the earliest opportunity.

The tip area in the north is brownfield and organic contamination (including benzo(a)pyrene) has been identified in exceedance of the threshold values and Hydrock believes barrier pipe is required where water supply pipes are to be place within this material. However, confirmation should be sought from the water supply company at the earliest opportunity.

6.4 Precautions Against Ground Gases

6.4.1 Radon

Current advice based on the BR 211 Report states that full radon protection is required for new dwellings across the entire site.

6.4.2 Landfill Gases

For the tip area in the north, all six of the required monitoring visits have been completed in previous investigations. The ground gas readings and gas regime conceptual model are considered to be sufficiently rigorous to provide an assessment of the ground gas regime and the scope of protection measures in accordance with CIRIA Report C665 (Wilson *et al* (2007).

For Situation B the Made Ground associated with the tip area in the northern part of the site, may be classified as "Green" and no protection would be required in new build in respect of methane or carbon dioxide in near surface soils.

6.5 <u>Waste Management</u>

Any material excavated on site may be classified as waste and it is the responsibility of the holder of a material to form their own view on whether or not it is waste. This includes determining when waste that has been treated in some way can cease to be classed as waste for a particular purpose.

One of the ways this can be achieved is set out in the Development Industry Code of Practice (CoP) (CL:AIRE, March 2011). This builds on the Environment Agency guidance document *Definition of waste: developing greenfield and brownfield sites* (2006).

The handling, re-use or disposal of waste is regulated by the Environment Agency. The Agency will take into account the use of the CoP in deciding whether to regulate materials as waste. If materials are dealt with in accordance with the CoP, the Agency considers that

those materials are unlikely to be waste at the point when they are to be used for the purpose of land development. This may be because the materials were never discarded in the first place, or because they have been submitted to a recovery operation and have been completely recovered so that they have ceased to be waste.

Further details of the CoP and the classification of waste are presented in Appendix D.

6.6 Outline Remedial Strategy

A cut and fill remedial strategy for the site will have to be developed in consultation with the design team and the regulatory authorities. Liaison should be continued during implementation and subsequent verification. The following approach is suggested for the area of Made Ground associated with the tip in the northern part of the site.

It is understood that the proposed development ground levels are to be raised over the majority of in this area to facilitate development, but there may also be some removal of materials in parts.

The material removed, if geotechnical suitable could be used to raise levels under roads and hard standing, or under proposed public open space, as long as a suitable cover or contaminant pathway breakage is in place to prevent direct contact with the Made Ground by end users of the site.

Through an appropriate Materials Management Plan (e.g. reuse of Made Ground on site with appropriate capping system) and on site processing of materials (e.g. separation of inert materials) the development will keep as much of the Made Ground material on site as possible, thus minimising vehicle movements and landfilling.

The 'fine' fraction, containing the soil, will require sampling (e.g. one sample per 500m³) to aid in the assessment of its suitability for re-use on the site. Depending on the results of the testing, the soils generated from the screening process will either be reused on site as either clean cover or, if the material is contaminated, as fill material to raise the site levels, where required and be covered by an appropriate clean cover.

Areas of Made Ground under gardens and public open space will require a clean cover of a minimum of 600mm thickness (designed in accordance with BRE guidance (Hollingworth 2004)). The extent and total thickness will need to be estimated from knowledge of the existing and final ground levels.

Foundations will have to be taken down to natural ground and consideration given to protection of water supply pipes and other utilities.

Full radon protection measures will be required and these should be capable of preventing ingress of naphthalene vapours in the tip area.

The methodology for the remediation should be detailed in a Remediation Method Statement which will need to be submitted to the NHBC and the regulatory authorities for approval.

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Following completion of the above works verification reports, undertaken by a suitably qualified independent engineer will be required.

7.0 GEOTECHNICAL INTERPRETATION AND RECOMMENDATIONS

7.1 <u>Geotechnical Categorization of the Proposed Development</u>

Eurocode 7, Section 2 advocates the use of geotechnical categorization of the proposed structures to establish the design requirements. For the purposes of this investigation, the proposed structures have been classed as Geotechnical Category 1.

7.2 <u>Geotechnical Aspects of the Development</u>

Geotechnical aspects of the site are discussed in the following sections. Specific areas of potential concern are associated with the appropriate foundations and earthworks associated with the proposed development.

7.3 Site Preparation, Earthworks, Groundworks and Landscaping

A rising sewer main and water main cross the eastern edge of the site from north to south. During the site investigation, works were undertaken in an attempt to locate the rising main. A manhole on the north eastern boundary of the site was dewatered for inspection and though the sewer pipe was not seen, the base of the manhole was approximately 7.5 m bgl, indicating the significant depth of the rising main and thus explaining the lack of success in locating the rising main through hand digging south of this point.

Where the proposed development is in close proximity to existing live services, they must be located and a wayleave clearly marked prior to undertaking ground works in the vicinity. Excavation in close proximity to live services will need to ensure that the excavation walls are stable or appropriately battered to a safe angle.

Temporary slope stability works may be required in any deep excavations.

Whilst no buried obstructions were encountered in the wider site area, an obstruction consisting of a large boulder-sized section of brick wall was encountered within the Made Ground in the tip area. Obstructions like this may affect piling and it is recommended that an allowance be made for removal or breaking up of obstructions.

Following the removal of obstructions, excavations of shallow soils should be feasible for conventional plant and equipment.

Significant excavation into the Marlstone Rock Bed may require the use of pneumatic breakers or rock rippers and there will be a reduction in progress where their use is necessary.

Topsoil and unsuitable Made Ground should be removed from beneath all building and hard standing areas.

Spoil resulting from excavations within the Made Ground may be suitable for reuse as fill to raise site levels elsewhere within the tip area, subject to its suitability with respect to appropriate geotechnical and geo-environmental specifications. Contaminated material should not be re-used on any part of the site currently uncontaminated.

Excavations undertaken during site works in the eastern edge of the site encountered unstable conditions in the sand and given the loose nature of the shallow material associated with the Charmouth Mudstone Formation and the high water table in this area, all excavations in the eastern end of the site should be supported and groundwater control may also be needed. Well-pumping may be required for dewatering as sump pumping will increase the instability of unsupported ground in these conditions.

It is recommended that no site personnel enter any trenches unless there is adequate support and this has been assessed by a competent person.

Groundwater seepages were encountered in the trial pits at shallow depths elsewhere on the site. In places where the sides are stable, groundwater seepage during excavation can be dealt with by sump pumping.

Groundwater levels may vary from those at the time of the investigation, for example in response to seasonal fluctuations.

The proposed development aims to generally raise levels and where material needs to be removed for either geotechnical or geo-environmental reasons (such as installation of a suitable capping layer or the excavation of foundations trenches) it is considered that the reuse of existing soils as part of redevelopment proposals will be necessary. The earthworks will need to be undertaken under a Materials Management Plan to ensure the appropriate management and reuse of the existing soils. Where the earthworks perform a structural role, a suitable specification will be required.

7.4 Foundations

The recommendations in this report follow NHBC Standards Chapter 4.2 (2011).

The preliminary foundation designs in this section are based on the parameters given by the 2012 Wardell Armstrong Report in Section 2.9.8. Selection of geotechnical design parameters should be undertaken in conjunction with the design process and discussed in a separate Geotechnical Design Report. For the consideration of foundations the site has been split into three separate areas as a result of the different ground conditions encountered. These areas are as follows and are highlighted on the Foundation Zonation Plan (Drawing C12702-G004) in Appendix A.

- Tip Area at the northern end of the site underlain by Made Ground of varying thickness over the Charmouth Mudstone Formation;
- The Northern Area Outside of the Tip Area in the northern part of the site, south of the tip area, underlain by the Charmouth Mudstone Formation; and
- The Southern Area of the site underlain by the Whitby Mudstone Formation and the Marlstone Rock Bed.

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7.4.1 Development Foundations for the Tip Area

The proposed finished levels in this area are generally to rise.

It is estimated that the combined existing Made Ground and proposed land-raise fill along the western edge of the site ranges from 0.5m in the north western corner to 2.8m in the south western corner. As a result, it is anticipated that trench fill spread foundations should be appropriate for the proposed row of plots along the western boundary of the tip area, to found in the underlying natural soils. This is shown on the Foundation Zonation Plan (Drawing C12702-G004), but will require confirmation by way of a plot-by-plot foundation assessment. The Wardell Armstrong Report outlined that the clays in the Charmouth Mudstone Formation are subject to moderate to high volume change potential, which should be taken into account in the separate Geotechnical Design Report.

The depth of foundations should be designed and the formations inspected by a geotechnical engineer. Any sub-formation materials deemed as unsuitable such as soft or loose zones should be excavated and replaced with well compacted suitable granular fill or lean mix concrete.

Foundation excavations should be protected from water and inclement weather including frost and any water should be removed by pumping from a sump in the base of the excavation.

Foundations which span founding materials of different stiffness should have mesh reinforcement placed top and bottom of the foundation.

The estimated thickness of existing Made Ground and land-raise fill across the remainder of the tip area is approximately 4.3m in the centre (at TP7), falling to 2.5m on the eastern edge of the tip (at HTP06). In addition, there are unstable natural ground conditions along the eastern edge of the tip area at the bottom of the slope next to the Oxford Canal (caused by water seepage promoting running sand conditions). At this stage, piled foundations are recommended in these areas, as highlighted on the Foundation Zonation Plan (Drawing C12702-G004).

The piles should be designed to penetrate into the stiff clay of the Charmouth Mudstone, gaining support from shaft adhesion and end bearing in the Charmouth Mudstone. Negative skin friction should be allowed for in the Made Ground and land-raise fill.

Bored or continuous flight auger cast *in situ* concrete piles or driven preformed concrete piles are considered suitable for the ground conditions at the site. Care should be taken with cast *in situ* piles where collapse of the pile shaft or running sand conditions could lead to 'necking' of the pile. A specialist piling contractor should be consulted for selection of appropriate piling method. An assessment should be made if driven piles are considered, as these may be subject to excessive noise or vibration which may be unacceptable to neighbouring properties.

Piles should be installed through the Made Ground and the sand deposits into the Charmouth Mudstone to estimated depths of between 5 and 10 m, but the final design will depend on detailed assessment by the piling contractor.

A working platform will be required to be designed and constructed prior to arrival on site of tracked piling plant. This should be designed and installed in accordance with BR470 (BRE 2004) based on data on the piling plant in accordance with an FPS certificate for the rig loadings.

7.4.2 Development Foundations for the Northern Area Outside of the Tip Area

The proposed finished levels in the northern end of the site outside of the former tip are to be raised by up to 1m from the existing site levels. Therefore, if founding strata are found to be deeper than about 2m bgl then spread foundations would be inappropriate and a piling solution would be required.

A review of the historic exploratory holes in the area show the depth to competent founding strata ranges from 0.2 to 3m bgl with no obvious spatial pattern.

As a result of the variable ground conditions, a combination of spread foundations and piled foundations would be appropriate in this area. This would need to be assessed by a plot by plot basis.

Both the spread and pile foundation solutions should be designed to penetrate into the stiff clay of the Charmouth Mudstone.

The clays in the Charmouth Mudstone Formation are subject to moderate to high volume change potential these issues should be taken into account in the separate Geotechnical Design Report.

The depth of spread foundations should be designed and the formations inspected by a geotechnical engineer. Any sub-formation materials deemed as unsuitable such as soft or loose zones should be excavated and replaced with well compacted suitable granular fill or lean mix concrete.

Spread foundation excavations should be protected from water and inclement weather including frost and any water should be removed by pumping from a sump in the base of the excavation.

Spread foundations which span founding materials of different stiffness should have mesh reinforcement placed top and bottom of the foundation.

7.4.3 Development Foundations for the Southern Area of the Site

The proposed finished levels in the southern part of the site are to be raised by up to approximately 0.15m from the existing levels.

The Wardell Armstrong Report 2012 states that as strip foundations up to 1.0m depth below existing ground levels with a minimum a safe bearing pressure of 100kPa would be considered appropriate for properties within the south of the site. Depending upon local conditions and the type of building and loadings proposed higher bearing pressures may be appropriate. Localised softer zones may dictate the requirement for deepened foundations and or excavation / removal / controlled replacement operations to be carried out.

In response to the ground conditions that have been identified at the site, it is considered that a spread foundation solution is likely to be appropriate for the proposed development in the south of the site.

In accordance with NHBC Standards Chapter 4.2 a minimum foundation depth of 0.9m will be required for strip and trench fill foundations extending through any topsoil or Made Ground into the Marlstone Rock Bed low volume change potential clays.

The foundations should be placed within the firm to stiff clay of the Marlstone Rock Bed. Within these materials a safe bearing pressure of 125kPa can be assumed which includes a factor of safety of 3.0 against general shear failure and will limit total foundation settlement to less than 25mm for foundation widths up to 1m.

On the basis of the site investigation, founding strata is anticipated to be at between 0.25m bgl and 0.4m bgl. However, this is subject to confirmation by a suitable competent person during excavation.

The depth of foundations should be designed and the formations inspected by a geotechnical engineer. Any sub-formation materials deemed as unsuitable such as soft or loose zones should be excavated and replaced with well compacted suitable granular fill or lean mix concrete.

Foundation excavations should be protected from water and inclement weather including frost and any water should be removed by pumping from a sump in the base of the excavation.

Foundations which span founding materials of different stiffness should have mesh reinforcement placed top and bottom of the foundation.

7.5 Foundations and Plants

Deepening of spread foundations in accordance with NHBC Standards will be required where foundations are within the zone of influence of existing or proposed trees and proposed shrub planting in accordance with the measured soil shrinkage on each clay soil type. Where foundations are within the influence of trees and are deeper than 1.5m bgl, a suitable compressible material or void former will be required.

Where foundations require deepening to greater than 2.5m below ground level, they must be designed by an engineer, as specified in NHBC Technical Requirement R5.

7.6 Ground Floor Slabs

In the northern part of the site associated with the tip area as Made Ground is greater than 600mm thick and clay soils of moderate to high volume change potential are present at the site, it is recommended that suspended floor slabs should be adopted, in accordance with NHBC Standards.

Ground bearing slabs may be used in the south of the site if the following criteria are satisfied:

- the foundation depth (such as due to the influence of trees) is less than 1.5m;
- any fill is suitable, well-compacted granular material and less than 600mm thick; and
- it is demonstrated that desiccation is not present and soils are at their equilibrium moisture content.

Prior to the placement of the founding materials and the construction of the ground bearing floor slab, the sub-formation and formation will need to be inspected and checked by a geotechnical engineer to ensure the ground conditions are as expected. This is likely to include sufficient appropriate testing, carried out in accordance with the DMRB IAN 73/06, to confirm the ground conditions at time of construction are consistent with the previous design parameters derived from this ground investigation.

If low bearing and soft strata are identified at the formation, this should be reported to the Geotechnical Engineer immediately and remedial actions agreed.

Ground floor slabs should be designed to incorporate any gas mitigation measures that may be required as discussed within the previous sections of this report.

7.7 Roads and Pavements

As part of the Wardell Armstrong 2012 report a total of ten CBR tests were undertaken across the site area generally within the locations of the proposed roads to be installed as part of the development works. The results of the testing indicate that CBR results at depths of *circa* 0.5m below existing ground level may be expected to vary from 3.0% to 12.0% within the southern section of the site close to Oxford Road (CBR104 to CBR110) with CBR values of between 0.9% and 1.6% within the smaller development area in the northern section of the site adjacent to the Oxford Canal.

The results of the CBR testing may be used to inform the detailed pavement design and appropriate sub-grade thicknesses.

Based on the test results, it is considered likely a CBR of 5% will be achievable over the south of the site of the site and can be used for preliminary design, subject to *in situ* testing during construction.

Proof rolling of the formation level will be required and any loose or soft spots to be removed and replaced with an engineered fill, in accordance with a suitable Specification. The formation level will also need to be protected during inclement weather from deterioration; all slopes shall be trimmed to falls to shed rain water and the surface sealed to limit infiltration.

Prior to the placement of the founding materials and the construction of the road pavement, the sub-formation and formation will need to be inspected and checked in accordance with a suitable Specification to ensure the ground conditions are as expected. All testing should be carried out in accordance with DMRB IAN 73/06 and confirm that the ground conditions at time of construction are consistent with the previous design parameters.

In the northern part of the site where the CBR is found to be less than 2.5%, the sub-grade may be unsuitable for both the trafficking of site plant and as support for a permanent foundation, without improvement works being undertaken. Improvement works should be

carried out in accordance with DMRB IAN 73/06 Rev 1 Chapter 5. In summary, consideration may be given to the following potential remedial techniques:

- excavation and re-engineering or replacement of weaker soils;
- the inclusion of geosynthetic reinforcement within the unbound layers of the capping and sub-grade;
- where cohesive soils are present and they are deemed suitable for treatment with hydraulic binders, to employ modification and/or stabilisation techniques on the formation; and
- where granular soils are present, de-watering and re-engineering the formation.

7.8 Soakaways and Drainage

The 2012 Wardell Armstrong highlights that limited soakaway testing was undertaken at the site but details are not available and it is possible that further soakaway testing may be required at the site.

7.9 Buried Concrete

Wardell Armstrong report 2012 undertook testing to assess the nature of the materials at the site in relation to the BRE Special Digest SD-1 – Concrete in Aggressive Ground. Twenty samples of soils were scheduled for geochemical testing to determine the sulphate content and pH of the materials at the site. The results were assessed in accordance with the methodology set out within the BRE Special Digest SD-1. The results indicate that the materials at the site would be classified as Design Sulfate Class DS-1 and Aggressive Chemical Environment for Concrete (ACEC) Class AC-1d.

7.10 Interaction Between Geotechnical and Geo-environmental Recommendations

An integrated approach to geo-environmental and geotechnical design solutions is required in order to derive the best option for site development.

From a geo-environmental perspective the tip area in the northern end of the site will require a 600 mm capping layer. The ground level is also to be raised to produce suitable final levels. The imported fill should be designed to serve both of these purposes. In order to ensure a minimum cover of 600 mm, some re-profiling of the Made Ground will be necessary.

Where piles are required, a risk assessment will have to be undertaken to demonstrate that no new contaminant migration pathways will be created by piling.

8.0 UNCERTAINTIES AND LIMITATIONS

8.1 General Comments

This report details the findings of work carried out in November 2012 and March 2013. The report has been prepared by Hydrock on the basis of available information obtained during the study period. Although every reasonable effort has been made to gather all relevant information, all potential environmental constraints or liabilities associated with the site may not have been revealed.

The report has been prepared for the exclusive benefit of Bovis Homes, Barratt Homes and Taylor Wimpey Homes and those parties designated by them for the purpose of providing geotechnical and geo-environmental recommendations for the site. The report contents should only be used in that context. Furthermore, new information, changed practices or new legislation may necessitate revised interpretation of the report after the date of its submission.

Hydrock has used reasonable skill, care and diligence in the design of the investigation of the site. The inherent variation of ground conditions allows only definition of the actual conditions at the locations and depths of trial pits and boreholes at the time of the investigation. At intermediate locations, conditions can only be inferred.

Groundwater findings described are only representative of the dates on which they were made and levels may vary.

Information provided by third parties has been used in good faith and is taken at face value; however, Hydrock cannot guarantee its accuracy or completeness.

The work has been carried out in general accordance with recognised best practice as detailed in guidance documents such as the CLR 11 Model Procedures (Environment Agency 2004), BS5930:1999 +A2:2010 and BS10175:2011. Important aspects of the risk assessment process are transparency and justification. The rationale behind the assessments carried out for this report is given in Appendix D. Unless otherwise stated, no assessment has been made for the presence of radioactive substances or unexploded ordnance. Where the phrase "suitable for use" is used in this report, it is in keeping with the terminology used in planning control and does not imply any specific warranty or guarantee offered by Hydrock.

The chemical analyses reported were scheduled for the purposes of risk assessment with respect to human health, plant life, ecosystems and controlled waters as discussed in the report. Whilst the results may be useful in applying the Hazardous Waste Assessment Methodology given in Environment Agency Technical Guidance WM2, they are not primarily intended for that purpose and additional analysis may be required should waste classification be required for consideration of off-site disposal of contaminated soils. Separate analyses will be required to meet the Waste Acceptance Criteria for specific landfill sites.

Unless otherwise stated, the chemical testing carried out for this report was not scoped to comply with the requirements of the water supply company and further work may be required.

The preliminary risk assessment process may identify potential risks to site demolition and redevelopment workers. However, consideration of occupational health and safety issues is beyond the scope of this report.

Please note that notwithstanding any site observations concerning the presence or otherwise of archaeological sites, asbestos-containing materials or invasive weeds such as Japanese knotweed, this report does not constitute a formal survey of these potential hazards.

Any site boundary line depicted on plans does not imply legal ownership of land.

8.2 <u>Site-Specific Comments</u>

Risks to human health from naturally occurring vanadium is not considered significant, but at the time of writing this has yet to be confirmed with the Local Authority.

Asbestos-containing materials were identified in a single sample from the tip area during one of the previous investigations. None was encountered by Hydrock, but the possibility remains that these materials may be present in the Made Ground.

Risks to the surface water environment are based mainly on eluate chemistry and although the EQS for copper and PAHs are exceeded in the Made Ground, the conceptual site model does not suggest a significant risk. Similarly, exceedances of the EQS for metals and PAHs from the natural soils are not considered a significant risk and would be no different from other areas of similar natural geology. These assessments should be confirmed with the Environment Agency. A piling risk assessment is likely to be required in the tip area to demonstrate the lack of establishing any new contaminant pathway.

The recommended use of standard water supply pipe over most of the site and barrier pipe within the Made Ground will require confirmation from the water supply company.

There is the potential for running sand conditions to develop in excavations close to the canal where silty or sandy soils are combined with a high water table. In these conditions it may be necessary to employ well-point dewatering before stable excavations can be formed.

The practical cut-off between trenchfill and piled foundations is normally taken to be a founding depth of 2-2.5 mbgl. It has been shown that potential foundation depths exceed this threshold in the northern part of the site. The areas where piling is likely to be required have been highlighted, but this is a provisional finding and more detailed assessment will be needed on a ploy by plot basis.

Existing CBR testing is sufficient for preliminary design, but confirmatory testing will be required in actual road foundation locations.

The 2012 Wardell Armstrong highlights that limited soakaway testing was undertaken at the site but details are not available and it is possible that further soakaway testing may be required at the site.

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9.0 RECOMMENDATIONS FOR FURTHER WORK

- Liaison with the NHBC, Contaminated Land Officer and Environment Agency concerning the findings of this report in respect of contamination.
- Confirmation with the water company with respect to water supply pipe materials.
- Provision of a piling contamination risk assessment.
- Provision of a Geotechnical Design Report including design and possible further geotechnical investigation for (a) spread footings (including suitability of plots, safe bearing pressure and mitigation of the effects of trees and shrubs); and (b) piled foundations (including specialist sub-contractor design and design of a piling mat for tracked plant).
- Soakaway testing in areas where these may be feasible, ie granular soils without high water table.
- Provision of a Materials Management Plan for re-use of site arisings.
- Provision of a Remedial Method Statement for the cover over the tip area and agreement with the regulatory authorities; and
- Verification of the remedial works on completion.

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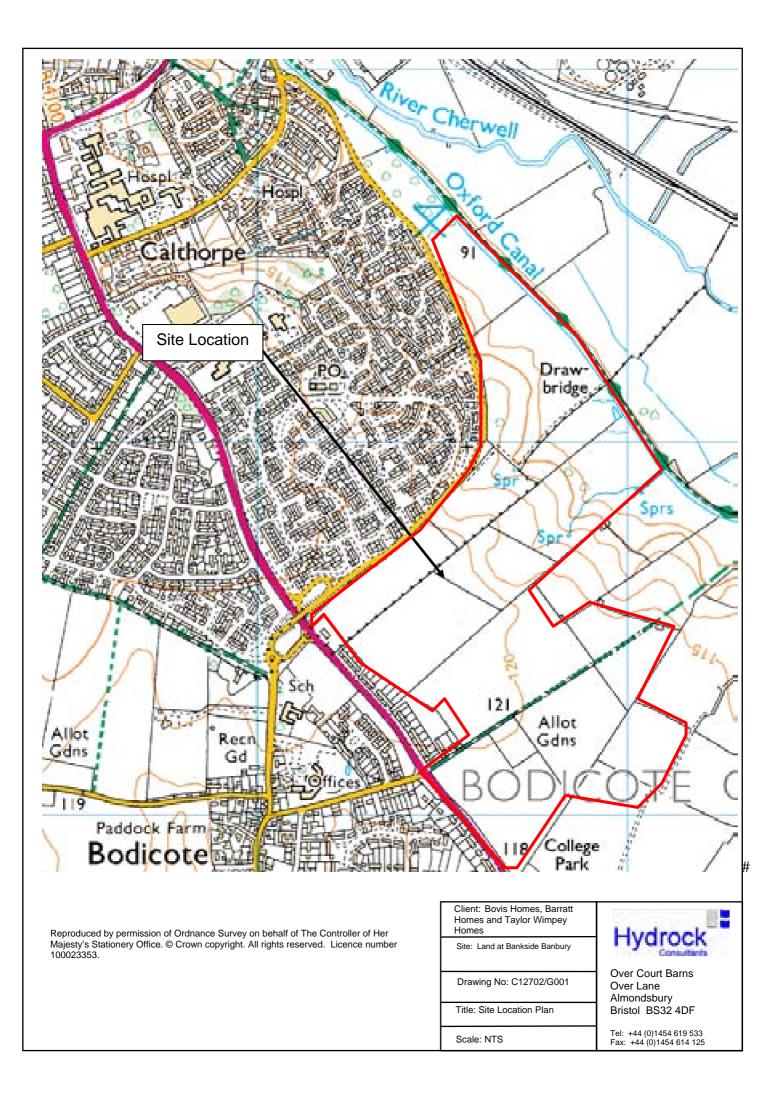
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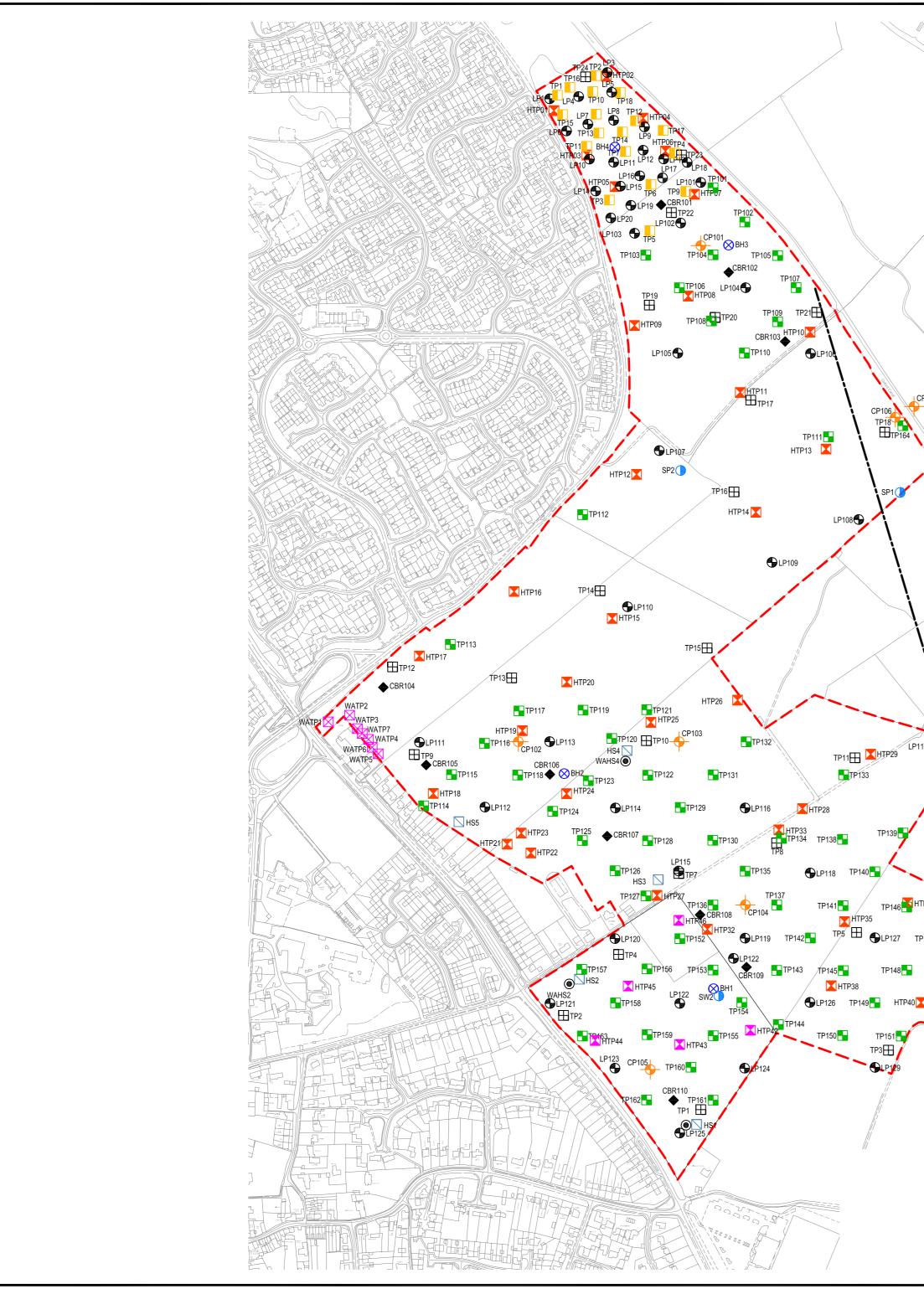
Appendix A

DRAWINGS

Drawings Included in this report:

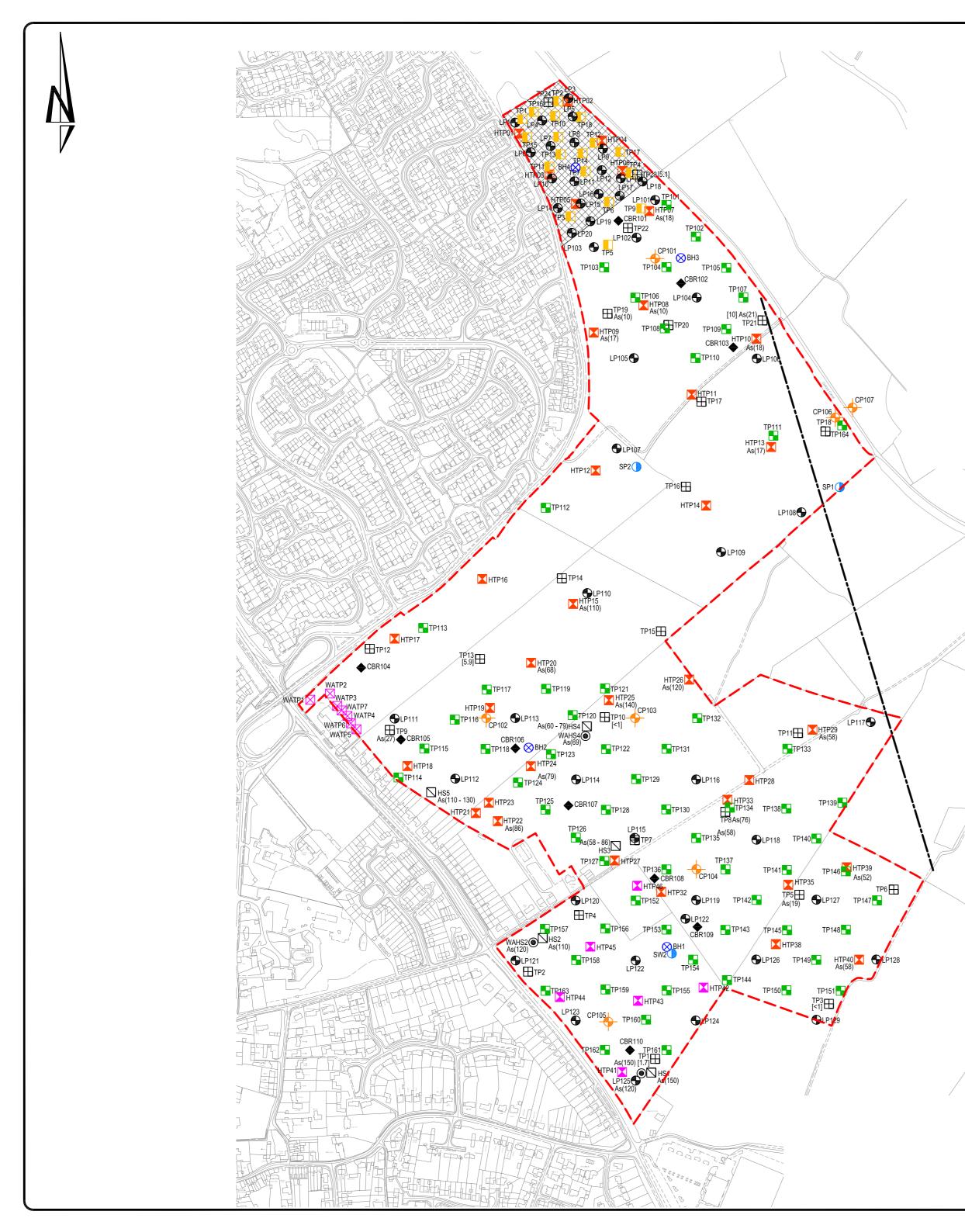
- C12702/G001 Site Location Plan
- C12702/G002B Exploratory Hole Location Plan
- C12702/G003B Geo-environmental Zonation Plan
- C12702/G004 Foundation Zonation Plan





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	HYDROCK APRIL 2013 TRIAL PIT SITE BOUNDARY	<u>No</u>	<u>tes:</u>					
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_	HISTORICAL WELL							
	HISTORICAL TRIAL PIT - 2012							
	HISTORICAL TRIAL PIT - APRIL 2006							
	HISTORICAL TRIAL PIT - 2005							
	HISTORICAL TRIAL PIT - MAY 2007							
	HISTORICAL HAND SAMPLE - JUNE 2007							
	HISTORICAL TRIAL PIT - DEC 2011							
⊗	HISTORICAL BOREHOLE - APRIL 2006							
•	CABLE PERCUSSION BOREHOLE							
	CBR TEST							
	WATER SAMPLE							
	RISING MAIN (ASSUMED ROUTE)							
As(21)	ARSENIC (mg/kg)							
[5.1]	PBET (%)							
	TIP AREA							
	GEOLOGY							
	GEOLOGICAL BOUNDARY							
	WHITBY MUDSTONE FORMATION (MAINLY CLAY)							
	MARLSTONE ROCK BED	В	05.04.13		Trial Pi	ts Added	NG	RS
	DYRHAM FORMATION (CLAYS & SILTS)	A	18/01/13		Issued for	Information	NG	RS
	CHARMOUTH MUDSTONE FORMATION (MAINLY CLAY)	Rev	Date		Desc	cription	Ву	Ckd
	NOTE - GEOLOGY BASED ON WARDELL ARMSTRONG 2012 REMEDIATION GROUND INVESTIGATION REPORT.	O` O` AI Br TE F <i>I</i> E- or CI		Barns 4DF 619 533 614 125 bl@hydro hydrock.o 6 HON	ck.com com MES, BA			6
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	specifications. This drawing is copyright.			C127	702 - G0	03	B	



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			specifications. This drawing is copyright.							1	· · ·]

Appendix B

SITE PHOTOGRAPHS

This appendix may not be included in the printed report to reduce the document size. It is presented in the PDF version of the report on the CD enclosed with the printed report.



Figure 1: Showing arisings from HTP01.



Figure 2: Showing arisings from HTP01.



Figure 3: Showing arisings from HTP01.



Figure 4: Showing arisings from HTP01.



Figure 5: Showing HTP01.



Figure 6: Showing arisings from HTP02.



Figure 7: Showing arisings from HTP02.



Figure 8: Showing HTP02.



Figure 9: Showing arisings from HTP03.



Figure 10: Showing arisings from HTP03.



Figure 11: Showing HTP03.



Figure 12: Showing arisings from HTP04.



Figure 13: Showing arisings from HTP04.



Figure 14: Showing HTP04.



Figure 15: Showing arisings from HTP05.



Figure 16: Showing arisings from HTP05.



Figure 17: Showing arisings from HTP05.



Figure 18: Showing HTP05.



Figure 19: Showing arisings from HTP06.



Figure 20: Showing arisings from HTP06.



Figure 21: Showing HTP06.



Figure 22: Showing arisings from HTP07.



Figure 23: Showing HTP07.



Figure 24: Showing arisings from HTP08.



Figure 25: Showing arisings from HTP08.



Figure 26: Showing arisings from HTP08.



Figure 27: Showing HTP08.



Figure 28: Showing arisings from HTP09.



Figure 29: Showing arisings from HTP09.



Figure 30: Showing arisings from HTP09.



Figure 31: Showing HTP09.



Figure 32: Showing arisings from HTP10.



Figure 33: Showing arisings from HTP10.

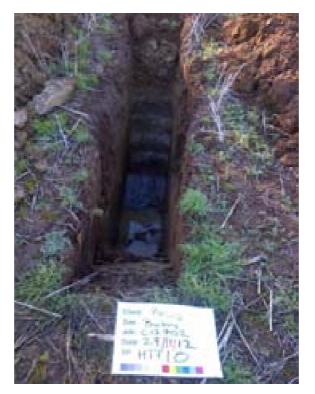


Figure 34: Showing HTP10.



Figure 35: Showing HTP10.



Figure 36: Showing arisings from HTP11.



Figure 37: Showing arisings from HTP11.



Figure 38: Showing arisings from HTP11.



Figure 39: Showing HTP11.



Figure 40: Showing arisings from HTP12.



Figure 41: Showing HTP12.



Figure 42: Showing arisings from HTP13.



Figure 43: Showing HTP13.



Figure 44: Showing arisings from HTP14.

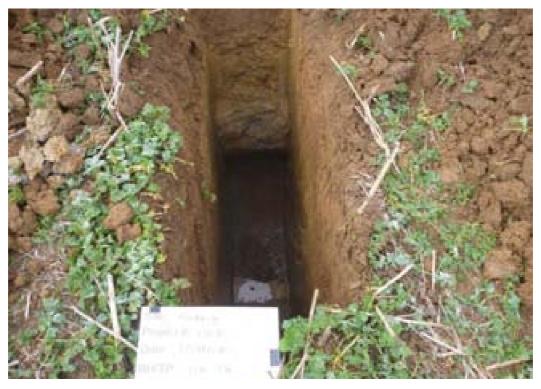


Figure 45: Showing HTP14.



Figure 46: Showing arisings from HTP15.



Figure 47: Showing HTP15.



Figure 48: Showing arisings from HTP16.



Figure 49: Showing HTP16.



Figure 50: Showing arisings from HTP17.



Figure 51: Showing HTP17.



Figure 52: Showing arisings from HTP18.



Figure 53: Showing HTP18.



Figure 54: Showing arisings from HTP19.



Figure 55: Showing HTP19.



Figure 56: Showing arisings from HTP20.



Figure 57: Showing HTP20.



Figure 58: Showing arisings from HTP21.



Figure 59: Showing HTP21.



Figure 60: Showing arisings from HTP22.



Figure 61: Showing HTP22.



Figure 62: Showing arisings from HTP23.



Figure 63: Showing arisings from HTP23.



Figure 64: Showing arisings from HTP24.



Figure 65: Showing HTP24.



Figure 66: Showing arisings from HTP25.



Figure 67: Showing HTP25.



Figure 68: Showing arisings from HTP26.



Figure 69: Showing HTP26.



Figure 70: Showing HTP27



Figure 71: Showing arisings from HTP28.



Figure 72: Showing HTP28.



Figure 73: Showing arisings from HTP29.



Figure 74: Showing HTP29.



Figure 75: Showing arisings from HTP32.



Figure 76: Showing HTP32.



Figure 77: Showing arisings from HTP33.



Figure 78: Showing HTP33.



Figure 79: Showing arisings from HTP35.



Figure 80: Showing HTP35.



Figure 81: Showing arisings from HTP38.



Figure 82: Showing HTP38.



Figure 83: Showing arisings from HTP39.



Figure 84: Showing HTP39.



Figure 85: Showing arisings from HTP40.



Figure 86: Showing HTP40.



Figure 87: Showing the Rising Main manhole at the northern end of the site adjacent to the canal.



Figure 88: Showing the Rising Main manhole at the northern end of the site adjacent to the canal (RM1).

Hydrock Consultants



Figure 89: Showing the 7.5m deep Rising Main manhole at the northern end of the site adjacent to the canal (RM1).



Figure 90: Showing the Rising Main manhole at the southern end of the site (RM3).



Figure 89: Showing the 7.5m deep Rising Main manhole at the northern end of the site adjacent to the canal (RM1).



Figure 90: Showing the Rising Main manhole at the southern end of the site (RM3).



Figure 91: Showing HTP41.



Figure 92: Showing arisings from HTP41.



Figure 93: Showing HTP42.



Figure 94: Showing arisings from HTP42.



Figure 95: Showing HTP43.



Figure 96: Showing arisings from HTP43.



Figure 97: Showing HTP44.



Figure 98: Showing arisings from HTP44.



Figure 99: Showing HTP45.



Figure 100: Showing arisings from HTP45.



Figure 101: Showing HTP46.



Figure 102: Showing arisings from HTP46.

Appendix C

EXPLORATORY HOLE LOGS

This appendix may not be included in the printed report to reduce the document size. It is presented in the PDF version of the report on the CD enclosed with the printed report.

Hydro		Dree Court flar Dree Lane Association Base Base Court Email Arout Tac Street Street Tac Street Street Tac Street S			Project: Land at B Banbury		P	Trial F HTI age N	PC	11 of		,
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				Very soft red brown line very sandy residual Ci	LAY.	
			2	(CHARMOUTH MUDSTONE)		4 1 문
						. 1 는 분왕
						10 A. 200
				Moderately packed light orange brown recidual occasional gravel. Gravel is medium angular di mudultore.	aft red britert	
				(CHARMOUTH MUDSTONE)		
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Date: 28/11/2012 Co-Ords: 446556.66.239403.00 Ellevation: 98.25mAOD Date troom clayer TOPSOL with roots (MADE GROUND) Soft and occasionally from between samdy graved scassional cotobles. Oravel and cotobles are a coarse bricks concerete trick fragments and pt (MADE GROUND) High strength still blue gray and yellow brown (CAY with the veha of table documpoont (gr (MADE GROUND) High strength still blue gray and yellow brown (CAY with shift blue gray and yellow brown (CAY concord)	Plant: JCB 3CX		in in in its second sec		
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Stratur Dark brown claywy 10PSOE, with rests. (MADE GROUND) Soit and occasionally firm brown samdy gravel socarise bricks concrete tricks forgenents and pt (MADE GROUND) High strength still blue gray and yellow brown GAY with this veins all black docomposed gri (MADE GROUND) High strength still blue gray and yellow brown GAY with this veins all black docomposed gri (MADE GROUND)	n Description		57 i	- 10	
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(MADE GROUND) Soft and occasionally firm brown samply grawn socarise bricks downed tolick higgmants and plu (MADE GROUND) High strength still blue gray and pelice brown GAY with this value of black decomposed ge (MADE GROUND) High strength still blue gray and pelice brown GAY with this value of black decomposed ge	notive firestowal ess.			- 10	
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0.10 0 (MADE GROUND) 1 m m m m 1 m m m m 0.30 0 (MADE GROUND) 1 m m m 1 m m m 0.30 0 (MADE GROUND) 1 m m m 1 m m m 120 0 (MADE GROUND) 1 m m m 1 m m m 120 0 (MADE GROUND) 1 m m m 1 m m 120 0 (MADE GROUND) 1 m m 1 m m 120 0 (MADE GROUND) 1 m m 1 m m 120 0 (MADE GROUND) 1 m m 1 m m 120 0 (m m (m m 1 m m 120 0 (m m (m m 1 m m 246 0 (m m (m m (m m 246 0 (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m (m m		a second second second second second	and a faith search the	Groundwale		Straturt	Description			31	10	18	ł
0.56 0 0.56 0 1.20 0 1.20 0 2.66 0 2.66 0 1.20 0 1.20 0 2.66 0 2.66 0 2.66 0 1.20 0 1.20 0 2.66 0 2.66 0	8.10	0			Dark (MAD	brown olayey TOPSOL. IE GROUNDI					10 (M)		*
1.20 D Image: Characterity packed descripte between and braves clayrey lime (SAND). Image: Characterity (Characterity packed descripte between and braves). Image: Image: Characterity (Characterity (Charac	8.50	D			to co bould	ana brick leagments concrete leagments a ler of brick wall laid on its side.	vel is angular medium nd a very large	1			-		
120 D 200 D. 240 D. End of triad pit due to triad pit due to triad pit obligate.				-32									88
2.00 D. 2.40 D. Evel of total pit due to total pit colleges. Evel of total pit due to total pit colleges.					Mode (CHA	nately packed orange brown and brown ch AMOUTH MUDSTONE)	ayey fire SAVD						
2.00 D. 2.40 D. End of trial pit due to trial pit colleges. End of trial pit due to trial pit colleges.	1.20	D											A STATE
2.46 D Evel of total pit due to thist pit collepse.	2.06	D.									1.4		
End of trial products that pic collegies.												***	
	2.40	0			Cod (244						
	10.000111 ⁻⁰⁰⁰				-								
General Remarks: Trial Pt was unstable. Water seepage at 0.9m bgl.			enten at 6 See H	ai									

Hydro	ock	One Coult Ba One Late Almostatory Brutit B103 40F 5 met troot/P Te: 3145 414	wheel are	Project: Land at Ba Banbury	ankside,		rial Pit HTP po No.	07	7	,
Method: Tria	i Pit			Date: 28/11/2012	Logged By: F	RS .	Chec			k
Client: Bovie	, Barratt a	nd Taylor V	Vimpy.	Co-Ords:446722.76,239342.00	Pit Stability: I	Unstable	Dime	nsk	D	
Hydrock Pro	ject No: C	12702		Elevation:91.27mAOD	Plant: JCB 3	сx	1.4	-0		1
San	ples and In 3	Situ Testing	Groundwater	Circle	m Description				1.5	1
Depth (m)	Type	Peoulte	1000000	Soft red brown fine sandy clay 10PSOIL.	to crosse group		$\rightarrow l^2$	T)	1.51	
				(TOPSOL)					-	128
0.50 0.50	NESY D	600%	V	Medium shangth soft to firm and brown slight CLAY (CHARMOUTH MUDISTONE)	y form startely respictual					
1.00	MỹV D	125kPa		High strength still ovange brown and gray mot (CHARBOUTH MUDISTONE)	Bed registual CLAY				-	
							1			
				Moderately packed onange brown line (SAND) (CHARMOUTH MUDISTORE)				-	-41.07	100
2.06	o			(CHWIMOUTH MUDSTONE)					-	100000
General Rema Inal Pit was unst		opage at 0.8m	ugi.							

Hydro	ock	Over Court Bar Over Larre Almonthibary Briatal BILLI KOF E-mail transfer	witted over	Project: Land at Ba Banbury		Trial Pit I HTPC Page No. 1	8	
Method: Tria	al Pit	Sectored links		Date: 29/11/2012	Logged By: RS	Check		h
Client: Bovis		rd Tavlor V	Vimpy	Co-Ords:446713.12,239185.00	the second s			ſ
Hydrock Pro				Elevation:99.87mAOD	Plant: JCB 3CX		R	2
	rples and In S		C. C. C. C.	ENAMOR 93'01 UNOD	Pieric 200 Jun		11	
Digiti (m)	Tge	Plends	Grundwater	Stratu	m Description	37	1	1
6.10	D			Brown clayey TOPSOL. (10PSOL)				X
	u .			High strength still orange brown and grey mot (CHARMOUTH MUDSTONE)	fel residual CLAV.	444	***	
0.50 0.50 0.50	NGV D B	9047a						
1.00 1.00	HSV D	(104Pa	SZ.				-	
				Firm to still source brown and grey motified re frequent line to medium entrometry weak Brow dark red lithurelicits at 1.7m logi pCH4/IMOU/TH MUDISTORE[]	idual CLAY with ficts with occasional		=1	
1.00	0			they have be		1	***	
General Remar Tsal Pt was stabl		ye ail tre byt						

Hydro	ock	Over Court Ren Over Laire Renordbury Bristol Bittal 40F E-mail eventiding Tex 1404 (1911)	dark own		Project: Land at B Banbury	ankside,	1.1	Trial F HTI ge No	P0	9	1	1
Method: Tria	al Pit			Date	29/11/2012	Logged By		Che	ciu	nd i	Βγ.,	k
Client: Bovi:	s, Barratt a	ind Taylor W	limpy	Co-C	vds:446630.30,239140.0	0 Pit Stability	: Stable	Dim	en			100
Hydrock Pro	ject No: C	12702		Eleva	tion: 106.08mAOD	Plant: JC8	3CX		-			
Sa	rples and In	Situ Testing	Groutballer		61	um Description				F	-2	1
Chipth (H)	Тури	Pesulta	00000	Sec.	conar In strength firm light tasken line sandy	615035		-	11	21	Ŕ	1
0.20 0.00	D HEV	604Pa 1204Pa		High s	OIL) tempth time to still orange brown and ndy readual CLAY getting tess sandy what FORMATIONS	prey motified skytche			1			
0.70 0.70	о В	100kPg										
100	D		X	10YR	etill brown dark red and grey mother e medium extremely weak lithorefict AM FORMATION()				+=			
				Fam 6 (0/199	ange brown and grey method residue AM FORWATION()	FOLMY.				+ 100		88
2.50	D			discounted in	own silly testidual CLAY with occasio- stated of very soft CLAY and occasio- t of gray CLAY with decongoosed the AMFCHMATION(of marking company time.			100			
6.09				Fishing		nelle.						
General Remo	diar (-			
		oage at 2.1-n byl										

Method: Trial PI Date: 23/11/2012 Logged By: R3 Checked By: Dimensions Client: Bovis: Barrati and Taylor Wimpy Co-Cvtd::466001.12.239129.00 PIS atability: Stable Dimensions Samples and In 56:: Training Bovis: Bo	Hydro	ock				Project: Land at Ba Banbury	nkside,		Pit No P10		2
Hydrock Project No: C12702 Elevation: 90.35mAOD Plant: USB 3CX ref Samples and In Site Training means Bevalues: 90.35mAOD Plant: USB 3CX ref 0 defining teel means Bevalues: 90.35mAOD Plant: USB 3CX ref 0 defining teel means Bevalues: 90.35mAOD Plant: USB 3CX ref 0 defining teel means Bevalues: 90.35mAOD Plant: USB 3CX ref 0 defining teel means Bevalues: 90.35mAOD Bevalues: 90.35mAOD ref 0 defining teel means flatter teel: 1000001 flatter teel: 1000001 ref 1.00 0 defining 0 defining flatter teel: 1000001 ref ref 1.00 0 defining teel: 1000001 exection and provided restand CAV 1.00 0 tee: 1000001 exection ref ref ref 1.00 0 exection ref ref ref ref 1.00 0 exection ref ref ref	Method: Tria	al Pit			Date	29/11/2012	Logged By: RS				X
Hydrock Project Nor. C12702 Elevation:90.35mACD Plant: JCB.3CX +	Client: Bovis	s, Barratt	and Taylor V	Vimpy	Co-C	rds:446901.12,239129.00	Pit Stability: Stal	ble Dir		10 March 10	1
Depth Int Tex Render Object Aller Statutus Deptropleon Ext 2 = 2	lydrock Pro	ject No: C	12702		Eleva	ition: 90.35mAOD	Plant: JCB 3CX		1		
Depinting Tape Reads Control to the first of the	Sar	nples and In	Situ Testing		1				. 1	_0	
0.50 0.00 HOV 0 90/24 Image: Imag	Organi (m)	Type .	Penda	1000	Lawrence -		Description		115	13	ł
1.00 HSV SDAPa 1.30 HSV HSV 2.00 D HSV 1.30 HSV HSV	0.50 0.50 0.50	HOV O B	904Pa		High II	rength firm to still orange boown and grey	rofiel veidual				
1.20 MSW 135MPs			90×Pa							1.0.0.0.0	
220 0 First to still grays and slightly sparses boost mutible resoluti CLAV and hoggest medium existently wattil Nicosholds. (CHARROCITH MUDSFORE)	-38	1.2.1	Mer in							1	
200 0 200 0	120		1304/%8						1	1 E	
200 0 200 0									11	1.5	
256 0 256 0										E	
256 0 256 0											
200 0 200 0									144	L.F	
200 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0											
encral Remarks:	-	12			CHAR	MOUTH MUDISTONE)	L.				
Intervales.	2.00	- M									
Incoral Pierrarks:									**		-90
Incoral Pierrarks:										E	23
incoral Rerundus											-
						Andread The Price	10.+				
Deneral Romarks. Inal Pit was stable. Field drain inflow at 0 tim bgl.											
Hall Pit was stable. Field drain inflow at 0 dm bgl.											
	ial Pit was stabi	e. Field drain	inflow at 0.9m bp	ś.							
Lapped in general accordance with NGNEE 1998. Incorporating Ameridment 1.							<u>1996 - 1997 - 1997 - 1997</u>				

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Method: Tri	al Pit	The Distriction		Date: 29/11/2012	Logged By: R	je No Che		of 1 d By:	1
		and Taylor V	Vimov	Co-Ords:446793.40.239036.00	Pit Stability: U			ions:	Contraction of the local division of the loc
Hydrock Pro				Elevation:94.66mAOD	Plant: JCB 3C		1.444	1. Star	h) I
	and the second se	Situ Testing						I.	0 7
Digiti (vi)	Type	Persuits	Groundwalter	in the constant of the second s	Description		37		1
				Firm slightly sity sky TOPSOL. (TOPSOL) Medium to high strength firm to still brown sligh	dy silly residual		+-00	***	12
				(CHAPMOUTH MUDISTONE)					
0.50	HSV	60kPa							100
0.50 0.50	8							-	
									腔
									124
1.00	HSV	85APia		Medium to high strength linn snange brown and	grey brown mottled	-	-		1.2.3
1.10	0			residual CLAY with occasional gravel. Gravel in dark red black mutatore. (CHARMOUTH MUDSTONE)	kine to medium				물물문
				Providence of the second					123
			- 52	Moderately to tightly packed light orange brown		_	1.48	10.0	1999
1.50	D			Gravel is fine to medium angular muditone and (CHARMOUTH MUDSTONE)					29.442 29.443
	105%			High strength still take grey residual CLAY with brown matter and occasional dark brown roted :	a slight orange		- 10	414	
2.00 2.00	HSV D	140kPa		brown motile and occasional dark brown rolled ((CHARMOUTH MUDSTONE)	part matter.	- 1			297
L De sinni								1	
									ES3
								18	2.2.2
								1.3	200
							1	- 54	
									223
									1.1.1
									5-8-8
							iù,	+ 4	
3.50	D.			Still dark grey residual CLAY with frequent mod- lithowich, country to be portrawing	un extensity weak		im a		
				(CHARMOUTH MUDISTONE)	10				

Hydro	ock	Brian Brian	which care		Project: Land at Bar Banbury	nkside,		rial F HTT 28 No	P1	2	
Method: Tria	u Pit	tie storenis		Date	: 30/11/2012	Logged By:		and the second second		d By:	k
Client: Bovie	s, Barratt a	and Taylor V	Vimpy	Co-C	ords:446633.43,238910.00	Pit Stability	Unstable	Dim	ens	ionsu	1
Hydrock Pro	Contraction Constraints			Elev	ation:104.55mAOD	Plant: JCB	эсх		****	1844	
Start Depth (m)	and the second second	Situ Testing	Grandeder		Stratum	Description			17		1
Pellin kul	Type	Pearls		Soft to	tion testion slightly sandy slightly gravely	day TOPSOL with		-+	81	25.5	120
6.25				aburd (TOP)	art rodela. IOBJ						182
0.29	~								**	-	28
				(DYR)	rength soft arange brown silly residual CE AM FORMATION	AK.					83
0.50	HEV	4049a		1							Ēŝ
											Es
										-	Ę
	20000										R
1.00	H9V.	35kPa						- 53			Ê
			w.						144		E.
1.30	HSV	154Pa		Very It (DYR)	w shangth very sait grey wet very sity re AM PORMATION)	ekual CLAY.					
	00										
1.50	в										胶
											63
											té
								*		-	1
											臣
											E
19.691	war-										
2.50 2.55	HSV D	7xPa									R
				Enter	tial pit due to trial pit collepse.				•		
				-010	the process of the proceeding the track of				1		1
								1			
Genoral Remark											
Dial Phywas unau	ible. Water se	Hoxpe at 1.2m by	p								

Hydro		Amonthibury Brucci BD22 407 E-mail braintithy Tar_21404 816.5	end on	Project: Land at B Banbury		Pa	HTP	ol		1	
Method: Tri:				Date: 30/11/2012	Logged By:						
		and Taylor W	/impy	Co-Ords:446925.66,238949.0	Dimension						
Hydrock Pro	and the second second second		_	Elevation:92.62mAOD	CX						
Sar Dept (m)	Types and In Type	Situ Testing Peruta	Grandeste	Stat	um Description		8	8.	19	1	
				Soft dark brown slightly sandy sitty clay TOP (TOPSOR)	SOL.		-			120	
6.20	D		2		7841. 7844			**			
0.40	HOV	16474		Low shength and yellow books ally residual (CHANMOUTH MUDISTONIC)	CLAX.						
0.60-0.70											
0.80	HIW 2	3(A/Fa									
							11				
140	D			Firm light brown and grey motified sandy silty occasional 0.1 to 0.2m Mich sand layers. Run to 1.6m big making pit very unstable. (CHA/IMCSUTH MUDISTICNE)	repdual CLAY with ving savd from 1.1m				C. A. W. T.		
2.00 2.10	D HSV	SSAPa		Vedum dranght limit brown and grey motiles residual CLAY. (CHARMOUTH MUDISTONE)	t slightly samte site						
				Modenstaty packed light brown sitty line SAV to medium gravel. (CHARMOUTH MUDISTORE)	D with accessoral line						
								474			
3.00	0						1.	L	-		
				End of bial pit due to truit pit collapse.	94184						
General Roma Trial Pit was une		oopage at 0.3++ b	g.								

ock	Bristyl Bristyl	ang Manaka sara	Project: Land at Ba Banbury	ankside,			P14				
al Pit	And service fields		Date: 30/11/2012	Logged By: /		and the second se		100 C 100	K		
s, Barratt a	and Taylor V	Vimpy	Co-Ords:446817.47,238851.00	Pit Stability:	Stable	Dim	ensia	d			
ject No: C	12702		Elevation: 104.61mAOD	and the second second second second	сx	1	-	-			
pies and in	Situ Testing	Generation	Contra Co	er Desertetter		-	. 1	1.0	1.2		
Type	Penda	2023		Sector Streams	_	-	182	<u>1</u> 34	1		
D			codels. (10PSOL)	POUL HIS SOUGHE				L			
HSV	25nPa		Low to medium strength soft becoming liem be CLAY (CHAVENOUTH MUDISTONE)	own sity residual							
HŞY	554Pa		High strength still orange brown and grey mult	fed silv residual			- 94	18-1			
HEV B	854%		CLAY becoming sandy before 2 Sm tigt. (CHARMOUTH MUDSTONE)								
HŞV D	70499										
HEV	1017a							1.18			
P		¥	End of Index								
						1					
os: a. Watter serves	içe at 2.8m tişi.										
	Al Pit A, Barratt I ject No: C pies and in Tase 0 HSV HSV B HSV 0 HSV 0 HSV 0 HSV	A Pit A Barratt and Taylor V ject No: C12702 pies and in Situ Testing Tipe Peads Pier Staffs Pier Staff	Al Pit A Barratt and Taylor Wimpy ject No: C12702 pies and in Situ Testing Type Results 0 JUSA HSV 256Pa HSV 556Pa HSV 556Pa HSV 856Pa D 756Pa D 756Pa S 756Pa S 756Pa S 756Pa S 756Pa S 756Pa S 756Pa	Al Pit Control and Parameter and the second and the	All Pit Longeological and transformation of the source of	Part II Part of the second point arm is the second point arm i	Pick Science of y Losses Page Net Page Net Science of y Page Net Science of	Victor Bits of Decision Page No. 1 or Environment Section 2010 Page No. 1 or Checked Section 2010 <td>Chr. No. 1 of 1 Page No. 1 of 1 NI PH Date: 30/11/2012 Logged By: AH Checked By Dimension A, Barratt and Taylor Wimpy Co-Ords: 446817.47,238851.00 Pit Stability: Stable Dimension year Montal and Taylor Wimpy Co-Ords: 446817.47,238851.00 Pit Stability: Stable Dimension year Montal Stav Testing Boundaries Statum Description Image 1 Image 1 10 Maximum Soid is firm formal signify sandy sity day IOHSOL sim stocker Image 1 Image 1</td>	Chr. No. 1 of 1 Page No. 1 of 1 NI PH Date: 30/11/2012 Logged By: AH Checked By Dimension A, Barratt and Taylor Wimpy Co-Ords: 446817.47,238851.00 Pit Stability: Stable Dimension year Montal and Taylor Wimpy Co-Ords: 446817.47,238851.00 Pit Stability: Stable Dimension year Montal Stav Testing Boundaries Statum Description Image 1 Image 1 10 Maximum Soid is firm formal signify sandy sity day IOHSOL sim stocker Image 1 Image 1		

ct No: C	Ind Taylor Wimpy 12702 Situ Testing Results	Elevation:	46595.91,238688.00 119.30mAOD Strator	Logged By: AH	Page N Chi Din	icki	nd 8	¥. ,	k
oct No: C les and in: Type D HSV	12702 Situ Testing	Elevation:	119.30mAOD Stratur	and the second second second second	Din		slor	Į	1000
les and in: Type D	Situ Testing	Solt to Sm da	Stratur	Plant: JCB 3CX		1.494	p di		
Type D HSV	Grave	Solt to Srm day abundant root		Elevation: 119.30mAOD Plant: JCB 3GX					
D		Solt to Srm day abundant root		n Description		6.	ł	-8	1
HSV		abundant roof	the state is a second of the second state of t			33	21	13	4
	3627		k red brown slightly sandy silty ets and rare limestone gravels.	dig 10-500 km					
	404Pa	Medium strang residual CLAY	th firm red brown slightly sandy Gravel is line to medium tub a	stightly gravely router intestone and					
HEY	55474	international							
32								1.2	
	Constraints and								
HSV	658.9%					1	1		
0						1		E	2.52
								1	
						2.0		***	E.
HOP	115479	MARLSTONE	on grey and brange brown motil ROCK BED)	ed residual CLAY.					
- 192	1.075								6
0									đ
						1			
								F	
						Ľ		Ē	
						ŧ.		E	
						-			ŝ
		Very self-pery	with some orange motiling very -	sity residual CLAY.				T	ġ
		Person (not both	manufacture and and			1		E	
								÷	-
		- Markey and Com	and have received therein bandeds to a	d formers plants	_	-			iii Ei
		 Darlinity weath 	Hed LINESTONE.	of Politica Politick		1040		/948	ą
		Astusal.	1554 ST.			F			
e					_			÷	
No grounde	water encountered								
	нду 0 нду 0	HSV 554% D 554% 0 1154% 0 1154%	HSV SSA% MARISTONE HSV SSA% MARISTONE D 115APs Montesting D Maristone Maristone Maristone Maristone Maristone Maristone Maristone Maristone	HEV 554% HARLSTONE HOCK BED) HEV 654% 0 HEV 1154% 1154	HSV 550°a HSV 1150°a HSV	HSV SSA's INVESTIGATION ROCK HED) HSV 65M's D HSV 65M's D HSV 65M's D HSV 65M's D HSV 115M's D H	HSV SSAPa HOCK BED) HSV BAPA D HSV BAPA D HSV BAPA D HSV BAPA D HSV BAPA D HSV BAPA D HSV BAPA D HSV BAPA D HSV BAPA D HSV BAPA HSV BAPA	HEV SSAPs ANALY SS	HSV BSUPs POWDPS (MARLETONE ROCK BED) IIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIII

Hydro	Hydrock				Banbury				Trial Pit No. HTP16 ge No. 1 of 1					
Method: Tria	d Die			Piste	29/11/2012	Looped By: A		e No.	10	11	1			
Client: Bovis		und Taulor I	Nimeu		xds:446444.55,238729.0			Check	nsic	oy. ms:				
Hydrock Pro			ampy	1000		Plant: JCB 30				-				
and the second second second	plos ind, G			EXEV	ation: 122.33mAOD	Piant: JUD 3	~		1		-			
Dagath (w)	1,04	Resits	Orserdente		Shah.	m Description		1	7)	13				
1	1-1-1	1		Figure 2	rown slightly sandy slightly gravelly sill art rooties.	y day 10PSOL with		-+17	T	T	19.00			
0.15	0			(TOP)				1	ľ	•				
0.29-0.45				Brown coarte (MAR	slightly samly clayey destructured GR tob angular intestone with abundant STONE ROCK BED;	AVES, Gravel is fire to cotoles.								
1.00	D							-						
				Moder seath (MAR)	stely strong line grained thirty bedded wed LMESTONE STONE ROCK BED)	red Brown shelly		1	•		15			
				1995	Enter hart	artiste"		- 1						
								the second s						
Seneral Remark Yal Pit was unsta		Seater encounts	red.											
					Lugged in a	proval accordance with BSS	E30 1989. in	and the second	Arrow	enint				

Hydro	ck	Deer Court Ra Over Lane Atmostationy Brazz Brazz Brazz Court Innovation Tec Innite Energy	without some		Project: Land at Ba Banbury	nkside,	HTP1 Page No. 1			.1 of 1 /		
Method: Trial	Pit			Date: 29/11/2012 Logged By			٨H	1 II. S 2 II. S 3	cked B	and the second s		
Client: Bovis,	Barratt an	d Taylor V	Vimpy	Co-0	Ords: 446298.59,238630.00	Pit Stability:	Stable	Dim	ension	1		
Hydrock Proj	ect No; C1	2702		Elev	ation: 123.49mAOD	Plant: JCB 3	CX			- C		
Sam	vies and in Si	tu Testing	Granter		Statut	Description		- 0	tal.	18 1		
Dageth (m)	Type	Persuits		Firm					3P2c	51 5		
0.15			(109	dark brown slightly sandy alightly gravely surt rootets. Gravel is time timestone and SOL) on to Noh strength firm to still broach grave				14	-83			
850	HSV	63474		abure	im to Nigh strength firm to stiff brown grav Gravel is medium to coarse suit angule ant cotities from 6 for to 0 firm togl LSTICNEL ROCK BEID)	Sincefore with				Environmentation of the second sec		
1.00	1457	130474								110		
2010										- 533		
1.20 D	D			weat	sawy strong line grained thirty bedded re ened LIMESTONE LISTONE FROM REDI	á brown sihelity				-		
					, Burgi ha Pro							
General Remark		der encounten	ed.									
					Loggest in gr	real advertarios with B	1000.000	Incorporat	ng Amerika			

t No: C1	the interests nd Taylor V 12702 itu Testing Pesuta		Date: 29/11/2012 Co-Ords: 446320.05,238418.00 Elevation: 122.09mAOD Stratum Oak brown slightly sandy sity day TOPSOL w and care gravels. (TOPSOL) High encept form to soft orange brown slightly a residuel CLAY. Guards of firm to course sub an inverted of LAY. Guards of firm to course sub an inverted of LAY. Guards of firm to course sub an inverted of the to course sub an	Logged By: AH Pit Stability: Stable Plant: JCB 3CX Description	1	By A
t No: C1 s and In S Tipe D HSV	12702 itu Testing Pesuta		Elevation: 122.09mAOD Stratum Oak brown slightly sandy sity day TOPSOL w and care gravels. (TOPSOL) High strength firm to still orange brown slightly is residual CLAY. Guards of line in correspondence	Plant: JCB 3CX Description		11
n and In S Type D	Pesula	Osuntaate	Elevation: 122.09mAOD Stratum Oak brown slightly sandy sity day TOPSOL w and care gravels. (TOPSOL) High strength firm to still orange brown slightly is residual CLAY. Guards of line in correspondence	Description	1	11
D MSW	Pesula	Osuntaate	Cark brown slightly sarely sity day TOPSOL at and care graves. (TOPSOL) High strength firm to still orange brown slightly a residual CLAY. Gravets of line in cores slightly a	8 - Fridden 19		II I
D			Cark brown slightly sarely sity day TOPSOL at and care graves. (TOPSOL) High strength firm to still orange brown slightly a residual CLAY. Gravets of line in cores slightly a	8 - Fridden 19		H I
HSV	504Pa		And rare growth. (10PSOR) High energit firm to still orange brown slightly a residual CLAY. Gaunta of line in correspond on	ih alumlari sotleta	1.1	
-15-	904Pa		episitum CLAY. Ocasely of line in charge suit an		1.00	
8			Temperature information	andy gravelly gular timestone		
					-	12.2
0						
			INVARIATION FROM BEDI	brown shally		
				1.11+	- 1	
					1	
					Lant Corr	
o groundwo	HIT PROUTERS	u.				
				0 Moderately strong live gained thinky booksed red membrand LifeSIONE (MARESIONE ROOK BED) Instants Instants Instants Instants	p grundweter encountered	D Moderately strong line paired thirly bodded red been shelly which is the strong line paired thirly bodded red been shelly line paired the strong line paired the

Hydro	Hydrock				Project: Land at Bankside, Banbury			Trial Pit No. HTP19 Page No. 1 of 1						
Method: Tri	al Pit	Tyr goethe same	141	Date	r: 29/11/2012	Logged By:		age re Che				b		
Client: Bovi		and Taylor V	Nimew		Ords:446457.42,238514.00		_		imensione.					
Hydrock Pro	a subscription of the second		a mapy	de la contrata de la		Plant: JCB :	1	1000		- 14				
the set of the set of the set of the set of	and the second state of the	Situ Testing	-	15 HON	ation: 122.22mAOD	Piant: 200	154A	-				-		
Depth (int)	Type	Peaks	Groundwater		Stratu	m Description			37	3.	18			
				Set 1	a liem brown slightly sandy silly-cliey TO rs and rate grantis of lime store.	\$504, with abundant			T.			r.		
9.29	D			(102	SOL)					-		62		
0.20						unovo ess		_	1.0		- *	25		
0.40	HSV	4(34 ¹⁰ 1)		Grav	an lo Nigh fam to still light brown gravel I is line to coarse sub angular linestone di below 12m ltgt. LISTONE ROCK BED)	y tesidual CLAY, becoming very								
0.00	HSV	125476								- 10		Ē		
1.00 D											3			
				-				_	14	- 1	**			
				weat the	rately shong line grained thirty bodded ered LIMESTONE LISTONE ROCK BEDI	an proses sussify					19			
				Partie	al. Buid hait	with the literature of the lit								
					1.0000000	MORTLE.								
								1						
General Roma	dis:													
Fial Pit was stat	sie. No ground	water encounter	ed											
					Lagged to	printal acceptorum of the	anter ocean	humping	ing A	-	ad I			

Hydro	ock	Over Court Be Over Lave Almontothary Brand BS24 40F E-mail Investig To 11454 2111		Banbury				Trial Pit No. HTP20 age No. 1 of 1								
Method: Tria	al Pit	for these and		Date: 30/11/2012	-	Logged By: A										
Client: Bovis	s. Barratt a	nd Taylor V	Vintov							nensiona						
Hydrock Pro	and the second	1.1.1.1.1.1.T. 1.1.1.1.1		والمتصادية الراجع فالألبا المتجهدات والا		Plant: JCB 30				1000						
and the second second	rples and in i		E	Elevation:121.64mAOD Plant: JCB 3CX -						1 1						
(Depth (H)	Tate	Pends	Groundwater		Shatun	Description			12	$\frac{1}{2}$						
0.15	D			Soft to live brown slightly grave roofiels. (TOPSOIL)	etty sity day 10	frebrute rile 308										
0.50	HSV	194Ag		Nedum strength fam is still o residual CLAY, Gravel is line t (MARLSTONE ROCK BED)	range tersen sligt In coarse suit ang	ty sandy graveity dar limestone.				***						
0.00 D				Moderately strong line grained weathered LBMESTONE (MARLISTONE ROCK BED)	therty bedded an	2 8 1 ⁰										
lenoral Remar								-								
nal Pé was stats	e. No provrdw	alter ericountere	0.													
					Logical in gen	est accederate with \$55	100.1008.0	a second and	i kie	in the second	1.					

Hydro	ock	Over Court Bar Over Lane Annochbary Broad BS32 40F X mail broad[In	etast ora	Project: Land at Banbury	Bankside,	Н	al Pit No. TP21 No. 1 of 1
Method: Tria	al Pit	heistenit		Date: 30/11/2012	Logged By:	and the second se	hecked By:
Client: Bovis		nd Taylor V	(imov	Co-Ords: 446433.90.238339			imension
Hydrock Pro	A CONTRACTOR OF			Elevation: 121.48mAOD	Plant: JCB 3	100 C	a atra
	rples and in t					246.5	
Depth Inti	Type	Pesula	Orsundwatter	1	atum Description		17 11
0.29	D HSV	854Pa		Soft to firm brown slightly sandy slightly s aturation evolutio. Gravel is firm to metho (TOPSOE) High strength still light yellow brown sligh GLAV becoming destructured very gravel (MARLSTONE ROCK BED)	m sub rounded limestone.		1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
0.40 0.40 1.10	D	130696					
				Moderately strong line grained thirty bad	ied red brown shelly		
				weathaned LIMESTONE (NAVESTONE ROCK BED) Petroat			
							<u>A.I.I.I.</u>
General Rema		aber encountere	d	Lagge	d r general accordance with t	125335 1996. From	pauling Amerimani 1.

Hydrock	Over Lane Over Lane Almondulary Brand Brand E. mart brandphate for the set 121		Project: Land at Ba Banbury		Trial I HT Page N	P2	2		
Method: Trial Pit	SC102-0314		Date: 30/11/2012	Logged By: AH	100		d B		ŀ
Client: Bovis, Barratt an	d Taylor Wir	nov (Co-Ords: 446470.59,238326.00	Pit Stability: Stable	Dim	iens	ion	e -	
Hydrock Project No: C1	The second second		Elevation:121.31mAOD	Plant: JCB 3CX			144	÷.	
Samples and In Si			sovalion, rensiningour	F HERE BOOM ORDER	_	1	11		27
Depth (m) Type	Pearls 0	ourdeeter.	Shatur	Description		11	20	18	ł
0.20 D 0.50 MSV 0.60 B	50x7x		Yelice brown and pay transfer diayey GRAVEL manage explain brick converse and rare add. (TOPSOE) High stempth still slightly gravely residual (CLA destructured very gravely below 6 8m byt. On the explaint interactions. MARLESTORE ROOK BED) MARLESTORE ROOK BED) MARLESTORE ROOK BED) Interaction of the Procession of the procession (CARLESTORE ROOK BED) Nature from the second	T becoming well is fine to coarse d brown shelly					
loneral filemarks:		_							
inal Pit was stable. No groundwas	e enduraded.			and according with \$25230, 1996	2.66				

Hydro	ock	Over Court Be Over Lane Annovability Briefs Briefs Ennel Driver(B			Project: Land at Ba Banbury	nkside,		Trial F HTI	2	3	
Method: Tria	ul Pit	Te induitie	da	Dat	2: 30/11/2012	Logged By:		age No Che		or 1 d By:	1
	and the second second	and Taylor V	Nimmy		Ords: 446455. 70, 238356.00	Pit Stability:				long	1
lydrock Pro			Constant.		ation: 121.71mAOD	Plant: JCB 3				100	1
- F	5	Situ Testing	-	END	WID01-121.71109000	T IN COL	nen.			E.	÷.,
Digiti (m)	Type	People	Ocerteite		Shatur	Description			R		
0.20	o			High befor	incom slightly gravelly slightly sandy sity fant roothets. SORJ strangth still yellow trown sity residual C + 0.40m log.	al navera					Noxod
0.40-0.50	HSV	954Pa			elige brown slightly sandy very gravelity n LISTOME ROCK BED;	esidual CLAY.					CALCULAR DE LA CALCOLOGICA DE
1.20	D			Mpde	namly strong line grained thirty bedded in word LMESTONE.	d brown shelly					L-RAMA
					Social Tra Pro-						
General Roma Irial Pt was stat	Second Second	water encounter	nd.		Loggetting	eneral accordance with t	10/16/10 19/18	Happen	ting Ar		

Hydro	ock	Arrondsbury Brood Bittle Kthr E-real broad	-		Project: Land at Ba Banbury	unkside,	1	Frial Pit No. HTP24 se No. 1 of 1				
Method: Tria	al Pit	14.0404.001		Date	x 28/11/2012	Logged By:	the second se	Checked By				
Client: Bovis	s, Barratt a	and Taylor V	Vimov	Co-C	Ords: 446525.56.238417.00	and the second second second second		Dimensiogs				
Hydrock Pro				a second	ation:121.79mAOD	Plant: JCB :	1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.	100				
3. The second	and the second second	Situ Testing	1	L. WY	alight r21.7 gm9000	1 10110. 01010	num -	18				
Depth (re)	Type	Pesats	Oroundheden	l	Skatu	m Description		IP . IS				
0.20 0.40 0.60 0.70 0.80 0.80 1.26	D HSV D HSV D HSV	45474 75474 65474		Wedse Store Stare	Meduan singhtly sandy slightly gravelly sitly clay TOPSOL. Wet with allowdam resultars. (TOPSOL) Meduan to traps strength lim to still pelcer brown and gray brown dightly sandy slightly gravelly sitly rescaled CLAY. Gravel is first to correct sub-argular timestore. Becoming very gravelly testor 1.20m og. (MARLSTONE ROCK BED) Meduately storeg fire grained red brown shelly partially seasthered (MARLSTONE ROCK BED) Meduately storeg fire grained red brown shelly partially seasthered (MARLSTONE ROCK BED) Meduately storeg fire grained red brown shelly partially seasthered (MARLSTONE ROCK BED) Meduately storeg fire grained red brown shelly partially seasthered (MARLSTONE ROCK BED)							
ioneral Remar M Pit was unes	States and see	dealer encounte	red.									

Hydro	ock	Over Court Ba Over Larre Atmontbluey Braind B122 40F E-mail braintile	yetradi carn	Project: Land at Ba Banbury	ankside,	Trial HT Page N	P2	5		/
Method: Triz	il Pit	Ter, Stribe and	60. 	Date: 28/11/2012	Logged By: AH	and the second second	io. i ecke	-		6
Client: Bovis		ind Taylor V	Vienov	Co-Ords: 446655.55,238527.00		-	nems		t di serie d	-
Hydrock Pro			in the		Plant: JC8 3CX			P		
	yect not: G			Elevation: 120.86mAOD	Plant JOB JOA			1		_
Chert into	Tex	Situ Festing Results	Groundwater	Stratu	m Description		12	1	Ŕ	
				First dark brown slightly samty slightly gravel	ly sity day TOPSOIL				"b2	惋
0.15	0			with abundant coohin. (TOPSION)						ŝ
0.00	HOV	SONPa		Medium strength firm to still red brown sandy	residual CLAY.				1.5	
0.42	0			(MARLSTONE ROCK BED)			1	-		23
0.50	HSV	70kPa 25kPa		1888-1779-1885-1777-1			-			
0.60	D			High strength still yellow brown gravely resid from to coarse sub-angular limesione and abu- flectoming very gravely testos 1 30m bgl. (MARLSTONE ROCK BED)	ndant coljddes.					
1.40	0			Moderately strong line to coarse grained blue	gray and red brown		-			12
			3	banded occasionally shelly weathered LIMES (MARLSTONE ROCK BED)	fone.		1	-	-	ίĽ,
				Petral.	w 190 m					
							1			
General Remai	da:									
Trial Pit was stab	w. wa grauna	water brookings	ed.	Kagged in	क्रामार्थ सरस्थित स्था विधियम् ११	01. stype	uing A	rande	upril 1	

Hydro	ock	Over Court Bar Over Lang Almonthery Brand BT20 40F E-mail Screenigh Lac 2HM 8103	wind out		Project: Land at Bar Banbury	nkside,	Pz	Trial P HTF Ige No	26	
Method: Tri	al Pit			Date	: 28/11/2012	Logged By:		Che	cked	8у:
Client: Bovi	s, Barratt (and Taylor V	Vimpy	Co-C	xds:446789.29,238562.00	Pit Stability:	Stable	Dim	ensic	Z
Hydrock Pro	oject No: C	12702		Elov	ation:118.89mAOD	Plant: JCB 3	CX			
and the second state of the second	riples and In		Grandwater		/ Rept. or	Description				18
Depth (vr)	Type	Pesuks	10252012	Martin	m strendth firm dach formers stimbly same	defile the flag			h	I.B
0.10	8 HSV	504Pa		105	OIL with occasional small tragments of bl hunders roothers.	a and white policy			**	
0.50 0.50-0.80 0.60	0 8 HGV	1005/Pa		sesida angula	inergin fern to still red brown slightly sand al CLAY with occasional gravel, Gaset is r lambour. Becoming gravely below 5.7 (310NE ROCK BED)	line to medicine sub-				State is a children and
				firite to cobbile	If arange brown very gravely destructure to arse sub angular forestore and mudate 510AE ROCK BEC)	e CLAY. Gravel is ne with albundant			-14	131125555555555555555555555555555555555
				Varia	eak thirty bedded griy partially weathened	M DSTORE with				-
2.50	D			0002334	nal sheki AM PORMATION)					
					Josia for Prat	E.+		1		
Joneral Florran	89. J	ater encountered	4							
					A second in second	nd accordence with M				

Hydro	ock	Over Court Ba Dver Lane Amondulury Brand B532 40F E-nat sheed	attack core		Project: Land at Ba Banbury		Trial F HT Page N	P2	7	1	
Method: Tri	al Pit	The state price		Date	0: 28/11/2012	Logged By: AH	Ch				k
	s, Barratt a	nd Taylor V	Mimov	-	Ords:446664.97,238260.00	Pit Stability: Stable	1000				
	oject No: C	the second second		-	ation: 120.78mAOD	Plant: JCB 3CX		-	15		
	mples and In S		13274	ALC: NO		1		1	F		
Daget (m)	Type	Pesulta	Grandem	F.,	Stratum	Description		R	10	Ŗ	ł
				Firm	brown slightly samly slightly gravely sity SOL1	day TOPSOIL					
0.20	D			1.00							
0.45	HSV	50kPe		i sight	um to high strength firm to still light brown by gravelity silly residual CLAY. Gravel is to be timestorie.	slightly sandy Ine to coarse suite				1	
0.60	D			- DAME	THY MUDSTONE FORMATION			1		÷	
9.70	HSV	90×Pa									
			2						12		
1.30	0										
				weat	rately strong line grained thinly bedded re wered LANESTONE. LISTONE POXX BED1	d brown shelly					남
				Pata					-		-
							*				
General Rem Trial Pe was sta	eka: ble. Field draini	nhow at 0.5m 8	*		Lagged in the	mend accordance with \$55550 Titl	8. iconor			- Internet	

Hydro		Over Coult By Over Lane Amendology Brianz BODI ADF E-mail torotol Ter Debis Stat	which says		Project: Land at Ba Banbury	nkside,	Pa	Trial F HTI ige No	P2	8	1
Method: Tria	al Pit			Dat	e: 29/11/2012	Logged By:		Che	cke	id E	A COMPANY OF THE OWNER
Client: Bovis	s, Barratt a	and Taylor V	Vimpy	Co-	Ords: 446889.31,238394.00	Pit Stability:	Stable	Dim	ens	ign	8
fydrock Pro	ject No: C	12702		Elev	ation: 119.69mAOD	Plant: JCB 3	CX				
and the second second second	mpiles and in		Groundwater		Stratur	Description			17	1	181
Depth (rej	Type	Pesalts	11.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1	Set	o lim brown alightly sandy slightly graved	Alter States			88	£1	57
0.20	o			100	dart rootes. 904.)				***	- 10	89
0.40	HSV	454Pg		10001	um to high strength from to still light brown all CLAY. Doshel to the to-medium sub an cosmicned califormium concretions. They MUESTONE FORMATION()	slightly gravetly sitty gular limentone					10.000
0.70 0.70 0.70-0.89	HSV D B	554Pa								- 14	CLUB CLUB
1.30	0			00479	still light brown gravelly residual CLAY. G e sub angular limestone with abundant co LSTONE ROOK BED)	gvel is fire to colori.				++	
				partie	nitely strong line grained thirty bedded in Ay weathered LIMESTONE. LISTONE ROOK BED) al	d tecsen shelly			14	***	** **
								and a second of the second			
leneral Remar ral Pit was stab	The second second	aler encountere	a		er Silve b	wal americana wit. B					and 6.

Hydro	ock	Drer Court Bar Drer Lane Amundsbury Briatit BESS 404	•		Project: Land at Ba Banbury	inkside,		HTF		100			
		E-mail temperity for mode and	and the second			4		ige No					
Method: Tria		_		Date	1: 29/11/2012	Logged By:		Che			 All the second se		
Client: Bovis	s, Barratt a	nd Taylor V	Vimpy	Co-C	Ords:446994.60,238478.00	Pit Stability:	Stable	Dime	ms	lony			
Hydrock Pro	vject No: C	12702		Elev	ation: 117.52mAOD	Plant: JC8 :	3CX	-					
Sar	rples and in 5	Stu Testing	Grandham		Frank	n Description				Į.	.8 I		
Dageth (m)	Тури	Persuits	1 2002021				_	-	81	h	19		
6.15 0.20	0 HSV	25494		to cou (10P	inengih soli brown samdy gravelly clay T esa sub angular limestone with abundar SOIL)	orsone, oraver is to crootens			1.00	++			
0.40-0.50	•			Acte to	p brown slightly sandy clayety destructur coarse sub angular limestone with abu LSTONE NOCK BED)	el GAWEL Gravel ndart cooper.					ELECTION OF		
0.80	D			Mode	analy strong line grained medium bodde	d eed brown				1	H.C.C.		
					ered LIMESTONE LISTONE ROCK BEDy				1.00		••田		
				Refus	d					-			
						545512							
								and the second second second second					
General Roma Trial Pt was stat	all horners	ater encounter	ed										

Hydro	ock	Over Lane Abrondshury Brussi BILU 407 5. rost brussiphy Tec Enter 615 10		Project: Land at Ba Banbury	inkside,	P	HTPS	32
Method: Tri	al Pit			Date: 28/11/2012	Logged By:	AH	Check	ed By
Client: Bovi	s, Barratt i	and Taylor W	impy	Co-Ords: 446742.63.238208.00	Pit Stability:	Stable	Dimen	
Hydrock Pri	oject No: C	12702		Elevation:120.10mAOD	Plant JC8 3	XDI		
Sa	riples and In	Situ Testing	Grundester		- I		L.	1
Cepilli (m)	Tape -	Pleasts		and the second	m Description		1	2-31
8.29 8.40 0.40 0.50 0.60	D HSV HSV	90479 100679		Fem light brown slightly sandy sitty day TOP (TOPSOL) Pight strength still yellow brown gravetly resid free to coolse sub-argular to sub-bounded Lin (MARLSTONE PROOK BED) High strength still brown grey brown and yells sandy gravetly residual CLAY. Gravet is fire to	al CLAY. Grand is estore		14	1 201115-2020-03230
1.00	D D	110474	V	sardy gravelly residual CLAY. Gravel is line to innerstone with occasional cobbles. (MARESTONE ROCK BED) Medium strong line to coarse graned with this blue gray calcoresist partially regiment UM	to medium besided			1 1 1 1 1 1 1 1 1 1
				(MAR, STONE ROOK BED) Refuted	#14 =			
Jeneral Rema Nai Pit was stab	878	ego at 1 2m bgi.			eneral accordances with 10			1.1.120

Hydro	ock	Over Court Ba Over Lane Atmostability Broad	**	Project: Land at Ba Banbury	nkside,	HTP:			
		BOAD HOP E-mail traterille Tax. Ethios. 2011	y Back Arm	1259-12120		age No. 1			4
Method: Tria	vi Pit			Date: 28/11/2012	Logged By: AH	Check			K
Client: Bovis	, Barratt a	nd Taylor V	Vimpy	Co-Ords: 446852.95,238361.00	Pit Stability: Stable	Dimen		2	5
Hydrock Pro	ject No: C1	12702		Elevation: 119.80mAOD	Plant: JCB 3CX	4.99	-		-
	ples and In S	Stu Testing	Grauthater	Chest or	Description		1	18	1
Depith (m)	Type	Resids	100000	Firm brown slightly savdy silty day TOPSOL w		. 8	ţh	1	
				(TOPSOL)	CT ACONSENT FOCUES				Ċ.
125	D								S
	1.1								RS.
6.40	HŞY	80692		High strength still yolice trown slightly sandy a with occasional decayed rooters. (WHITEY MUDISTONE FORMATION)	Ry residual CLAY				
0.80	HSV	75kPa							
1.00-1.10	0			Sall yellow brown slightly sandy gravelly residu fine to coarse sub angular to sub rounded time (MARLSTONE ROCK BED)	al CLAY, Gravel is done	-	Γ		
	1223			4			+ 14		
1.29	D						Ľ		
1.45	0					1	1.00		
111202				Medium strong fine to coarse grained with this t blue grey: coloareous partiality weathered LME abundarit shells. (MMRLSTONE ROCK 8ED) Results	o medum teddwif STCNE with		12		
				Read of Free Price	an an an				
General Rema Trial Pit was stak		aler ercounter	ect						
				Lagged in g	reul acordana wê 82000 1996	inconjunating	Ameri	dii a nt	1

Hydro	ock	Over Court Be Over Lane Amorbibury Brane Blate 407 E-mail present	which care	Project: Land at B Banbury	ankside,		Trial Pit HTP Ige No.	35	
Method: Tri	al Pit	Tel: Direkt Arts 5		Date: 28/11/2012	Logged By:				Brick
Client: Bovi		od Toulor V	Vienese				Dimer	tiskoe	1 miles
and the second	and the second second second		rimpy	Co-Ords:446954.21,238220.0	• • • • • • • • • • • • • • • • • • •		11.12	4	2
Hydrock Pro	A state of the state of the		-	Elevation: 119.26mAOD	Plant: JC8 :	RJR.	1	-	
Depth (ret)	riplies and In S Texe	Preute	Grandvater	Strat	um Description		5	23	19 1
				Soit to liem light brown silly clay TOPSOL (TOPSOL) Medium strength firm Rgit yellow brown silg CLAY with occasional angular limestore col	tdy sandy silly residual idea how 6.7m byl				-
0.50 0.60 0.60-0.70	HSV D B	704Pa		and raw increations colddies before 1.20m by partners MUDISTONE FORMATION				-	
1.10 1.16	NSV D	90475	v						
				Still yellow brown gravely residual CLAY. Gravel is line to coarse				-	
				sub angular to angular increations with abund (MARLSTONE ROCK BED)	et cubbles.		1.000		
				Medium strong line to coarse-grained red bri (MARLSTONE POOK BED)	we PONSTONE.			-	
				Parland.			- 21		
_									
General Rema	rka:								
Triat Pit was stab	na tratar bolga	φσ in 1.641 bỹ			general accordance with D				

Hydro	ock	Over Court Barris Over Laire Annochbory Broad BS22 45# E-sail broad(Hudrock core To: broad broad(Hudrock core To: broad broad(H)		Project: Land at Ba Banbury	Trial Pit No. HTP38 Page No. 1 of			1	1 1		
Method: Tria	il Pit		Dat	e: 29/11/2012	Logged By:	AH	Che			10 A 10	K
Client: Bovis	, Barratt i	and Taylor Wimpy	Co-	Ords: 446933.62,238121.00	Pit Stability	Stable	Dim	ene	sion	£.	
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Appendix D

HYDROCK METHODOLOGY

Hydrock Report Appendix on Hydrock Methodology, version 02 updated 21-12-12 applies to this report.

This appendix may not be included in the printed report to reduce the document size.

It is presented in the PDF version of the report on the CD enclosed with the printed report. Alternatively, it can be supplied on request by quoting the version number and date.

1.0 HYDROCK REPORT APPENDIX ON HYDROCK METHODOLOGY

This appendix provides additional background information on certain approaches and methods used by Hydrock Consultants Ltd in the preparation of this report.

Throughout the report the term 'geotechnical' is used to describe aspects relating to the physical nature of the site (such as foundation requirements) and the term 'geoenvironmental' is used to describe aspects relating to ground-related environmental issues (such as potential contamination). However, it should be appreciated that this is an integrated investigation and these two main aspects are inter-related. The geoenvironmental sections are written in broad agreement with BS 10175:2011.

The **first stage** of a two-staged investigation and assessment of a site is the Preliminary Investigation (BS 10175:2011), often referred to as the Phase 1 Study¹, comprising desk study and walk-over survey, which culminates in the Preliminary Risk Assessment. A preliminary conceptual site model (CSM) is developed. From this are identified any geotechnical and geo-environmental hazards and the qualitative degree of risk associated with them. From the geo-environmental perspective, the Hazard Identification process uses professional judgement to evaluate all the hazards in terms of **possible contaminant linkages** (of source-pathway-receptor). Possible contaminant linkages are potentially unacceptable risks in terms of the current contaminated land regime legal framework and require either remediation or further assessment. These are normally addressed via intrusive ground investigation and generic risk assessment.

The **second stage** is the Ground Investigation, Generic Risk Assessment and Geotechnical Interpretation. This represents the further assessment mentioned above. The Ground Investigation comprises field work and laboratory testing based on the findings of the Preliminary Risk Assessment, to reduce uncertainty in the geotechnical and geoenvironmental hazard identification. This may include the Exploratory, Main and Supplementary Investigations described in BS 10175:2011.

For the geotechnical aspects of the report, the general requirements of Eurocode 7 (BS EN 1997-2:2007) are to produce a Ground Investigation Report (GIR) which shall form part of the Geotechnical Design Report (GDR). The geotechnical section of this report is intended to fulfil the general requirements of the GIR as outlined in BS EN 1997-2, Section 6.

The GIR contains the factual information including geological features and relevant data, and a geotechnical evaluation of the information stating the assumptions made in the interpretation of the test results.

¹ Please note that it does not refer to a site development phase.

2.0 SITE INVESTIGATION INFORMATION

2.1 <u>Unexploded Ordnance</u>

Clients have a legal duty under the CDM 2007 Regulations to provide designers and contractors with project-specific health and safety information needed to indentify hazards and risks. This includes the possibility of unexploded ordnance (UXO) being encountered on the site. Further details are given in CIRIA report C681 (Stone *et al* 2009).

A non-UXO specialist screening exercise has been carried out for the site by considering (a) any evidence of UK defence activities on or near the site evident from the gathered desk study information and (b) the unexploded aerial delivered bomb (UXB) regional risk maps produced by Zetica. Other data sources are available, but as a first stage screening exercise the freely available Zetica maps have been used. The level of risk stated is that determined by Zetica, a company experience in the desk study, field investigation and clearance of UX0/UXB.

2.2 <u>Hydrogeology</u>

Under the Water Framework Directive the designations of principal and secondary aquifers is based on the Environment Agency interactive aquifer designation map. Where aquifers have been mapped, and they are capable of sustaining a yield of 10 m³/day or supplying 50 people on a continuous basis, the Environment Agency has designated a number of Groundwater Bodies to help manage water quality under the River Basin Management Plans. Groundwater bodies are defined based on their support for ecosystems as well as their capacity to supply drinking water. Note that some localised small aquifers capable of supporting the above supply may be too small to map and can be identified only by investigation.

Where an aquifer exists and it contains groundwater but is incapable of sustaining the above supply, the groundwater is not part of a Groundwater Body and is not considered a strategic resource. In which case the groundwater is not a receptor, but can be a pathway to other receptors by virtue of its ability to transport contaminants.

2.3 Geotechnical Testing

Derived values of geotechnical parameters and/or coefficients are obtained from test results, by theory, correlation or empiricism in line with BS EN 1997-2:2007, Section 1.6.

Where derived geotechnical parameters are to be used in designs in accordance with EC7, there are two further stages of interpretation that will be carried out by the geotechnical designer. The first of these is the selection of **characteristic values** for geotechnical parameters using the derived values and complemented by well-established experience as per EN BS 1997-1:2004, Section 2.4.5.2. The characteristic value is a cautious estimate of the value affecting the occurrence of the limit state. Consequently, any particular material

type may have more than one characteristic value for each parameter because there may be more than one limit state depending what is being designed.

The second stage is the selection of **design values** as per EN BS 1997-1:2004, Section 2.4.6.2. The design values is either derived from the characteristic value by applying the relevant partial factor or is assessed directly. Similarly, there can be several design values for the same material type.

In the event that geotechnical designs are include in this report, selection of the characteristic and design values is included. Otherwise, it is the duty of the geotechnical designer to determine these within a separate design report.

3.0 RISK ASSESSMENT RATIONALE

The work presented in this report has been carried out in general accordance with recognised best practice as detailed in guidance documents such as in the CLR 11 Model Procedures (Environment Agency 2004a), BS5930:1999+A2:2010 and BS10175:2011. Important aspects of the risk assessment process are transparency and justification. The particular rationale behind the risk assessments presented is given in this appendix.

A preliminary risk assessment is made of both geotechnical and geo-environmental hazards identified at the desk study stage and confirmed (or amended) at the ground investigation stage. In the case of geo-environmental hazards this is based on a simple matrix of probability of occurrence versus the consequence, as explained below, and is referred to as the **exposure model**. In the case of the geotechnical hazard identification, this is referred to as the **ground model**.

The geo-environmental risk assessment process proceeds to the next level, the generic risk assessment, in which actual contaminant concentrations are considered.

3.1 Preliminary Risk Assessment

In line with the CLR 11 Model Procedures (Environment Agency 2004a), the Preliminary Risk Assessment includes a geo-environmental Hazard Identification, which seeks to list all the suspected contaminant **sources**, the **receptors** that might be harmed by those sources and the **pathways** via which the sources might reach the receptors to cause the harm. The source-pathway-receptor concept is known as a contaminant linkage (formerly a pollutant linkage) and only when a linkage is complete is there any possibility of risk of harm arising.

The Hazard Identification process uses professional judgement to evaluate all the hazards in terms of **possible contaminant linkages**. Possible contaminant linkages are potentially unacceptable risks in terms of the current contaminated land regime legal framework and require either remediation or further assessment. These are normally addressed via intrusive ground investigation and the chemical analysis of soil and water samples.

Where no ground investigation has been carried out (i.e. in a desk study only report) there is greater uncertainty in the information available and so a geoenvironmental consequences and probability assessment is undertaken.

Some linkages may be identified which constitute a theoretical connection between a source and a receptor, but professional judgement shows them not to be possible for some reason. These are labelled 'no linkage' in the summary table and no further action is required. If a linkage is possible, a comparison is made of consequence against probability in general accordance with the guidance given in CIRIA Report C552 (Rudland *et al* 2001).

Classification of consequences and probability are given in CIRIA C552 Tables 6.3 and 6.4, respectively, but there are a number of inconsistencies in the original Table 6.3, in particular

relating to 'significant harm or significant possibility of significant harm' (SH/SPOSH). Consequently, the table has been updated by Hydrock in line with current practice and the revision presented in R&D Publication 66, Annex 4 (NHBC and Environment Agency. 2008, and is given in Table 3.1 below.

The basis of the classification is that 'severe' and 'medium' are likely to result in SH/SPOSH as defined by the EPA 1990, Part 2A, with 'severe' resulting in acute harm. 'Mild' lies below the level of SH/SPOSH but above the level of 'no harm' as implied by the relevant Generic assessment criterion (GAC, see below). Minor lies below the 'no harm' level.

	Classification of Consequences for Ge	o-environmental Risks
Classification	Definition	Examples
Severe	Concentration of contaminants is likely to (or is known from previous data to) exceed that indicative of unacceptable intake or contact. Highly elevated concentrations likely to result in "significant harm" to human health as defined by the EPA 1990, Part 2A, if exposure occurs. I.e. >>SH/SPOSH, concentrations are high enough to cause acute (short-term) effects.	Human health: short-term (acute) effects likely to result in significant harm. E.g. high conc. of cyanide on the surface of an informal recreational area. Significant harm to humans is defined as death, disease*, serious injury, genetic mutation, birth defects or the impairment of reproductive functions. Planting: complete and rapid die-back of
	Equivalent to EA Category 1 pollution incident including persistent and/or extensive effects on water quality; leading to closure of a potable abstraction point; major impact on amenity value or major damage to agriculture or commerce. Major damage to aquatic or other ecosystems, which is likely to result in a substantial adverse change in its functioning or harm to a species of	Iandscaped areas. Controlled waters: short-term pollution, e.g. major spillage into controlled water. Major fish kill in surface water from large spillage of contaminants from site. Highly elevated concentrations of List I and II substances present in groundwater close to small potable abstraction (high sensitivity).
	special interest that endangers the long-term maintenance of the population. Catastrophic damage to crops, buildings or property.	Buildings etc.: catastrophic damage, e.g. explosion causing collapse. (can also equate to immediate human health risk if buildings are occupied). Ecosystems: acute risk to a particular ecosystem
		or organism forming part of that ecosystem in a designated protected area, e.g. by contamination spillage. Damage to a protected area of international significance (e.g. Ramsar site). Site workers: risk assessment required to determine PPE and this may involve USEPA Level A, B or C protection.

Table 3.1: Classification of Consequences of Geo-environmental Risks

	Classification of Consequences for Ge	eo-environmental Risks
Classification	Definition	Examples
Medium	Concentration of contaminants is likely to (or is known from previous data to) exceed that indicative of unacceptable intake or contact. Elevated concentrations which could result in "significant harm" to human health as defined by the EPA 1990, Part 2A if exposure occurs. I.e. >SH/SPOSH. Equivalent to EA Category 2 pollution incident including significant effect on water quality; notification required to abstractors; reduction in amenity value or significant damage to agriculture or commerce. Significant damage to aquatic or other ecosystems, which may result in a substantial adverse change in its functioning or harm to a species of special interest that may endanger the long-term maintenance of the population. Significant damage to crops, buildings or property.	 Human health: long-term (chronic) effects likely to result in significant harm. E.g. high conc. of contaminants close to the surface of a development site. Significant harm to humans is defined as death, disease*, serious injury, genetic mutation, birth defects or the impairment of reproductive functions. Planting: stressed or dead plants in landscaped areas. Controlled waters: pollution of sensitive water resources, e.g. leaching into principal or secondary aquifers or rivers. Buildings etc.: damage renders unsafe to occupy e.g. foundation damage resulting in instability. Ingress of contaminants through plastic potable water pipes. Ecosystems: chronic death of species in a particular ecosystem in a designated protected area, e.g. by contamination spillage. Damage to a protected area of national significance (e.g. Site of Special Scientific Interest). Site workers: risk assessment required to determine PPE and this may involve USEPA Level B, C or D protection.
Mild	Concentration of contaminants is likely to (or is known from previous data to) exceed that indicative of no harm but not unacceptable intake or contact. Exposure to human health unlikely to lead to "significant harm". I.e. >SVG/GAC but <sh sposh.<br="">Equivalent to EA Category 3 pollution incident including minimal or short lived effect on water quality; marginal effect on amenity value, agriculture or commerce. Minor or short lived damage to aquatic or other ecosystems, which is unlikely to result in a substantial adverse change in its functioning or harm to a species of special interest that would endanger the long-term maintenance of the population. Minor damage to crops, buildings or property.</sh>	 Human health: harm but probably not significant harm unless particularly sensitive individual within the receptor group. May be aesthetic/olfactory impacts. Exposure could lead to slight short-term effects (e.g. mild skin rash). Planting: damage to plants in landscaped areas, e.g. stunted growth, discoloration. Controlled waters: pollution of non-sensitive water bodies e.g. leaching into non-classified groundwater or minor ditches. Buildings etc.: damage to sensitive buildings etc. Surface spalling of concrete. Ecosystems: minor change in a particular ecosystem in a designated protected area, but not significant harm. Damage to a locally important area. Site workers: risk assessment required to determine PPE and this may involve USEPA Level C or D protection.

	Classification of Consequences for Geo-environmental Risks											
Classification	Definition	Examples										
Minor	Concentration of contaminants is likely to (or is known from previous data to) be less than that indicative of no harm. No measurable effects on humans. I.e. <sgv gac.<br="">Equivalent to insubstantial pollution incident with no observed effect on water quality or ecosystems. Repairable effects of damage to buildings, structures and services.</sgv>	No measurable effects, but simple PPE required (USEPA Level D protection, i.e. overalls, boots, goggles, hard hat). The loss of plants in a landscaping scheme. Discoloration of concrete.										

CIRIA Table 6.4 is reproduced as Table 3.2 below. This provides an estimate of the probability that the event described by the contaminant linkage will occur. For example, the likelihood that pollution of groundwater will occur by leaching of metals into the aquifer.

Table 3.2: Classification of Probability of	Geo-environmental Risks
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	Classification of Probability of Geo-environmental Risks
Classification	Definition
High Likelihood	There is a contaminant linkage and an event that either appears very likely in the short term and almost inevitable over the long term, or there is evidence at the receptor of harm or pollution.
Likely	There is a contaminant linkage and all the elements are present and in the right place, which means that it is probable that an event will occur. Circumstances are such that an event in not inevitable, but possible in the short term and likely over the long term.
Low Likelihood	There is a contaminant linkage and circumstances are possible under which an event could occur. However, it is no means certain that even over a longer period such event could take place, and is less likely in the shorter term.
Unlikely	There is a contaminant linkage but circumstances are such that it is improbable that an event would occur even in the very long term.

The perceived level of risk for each pathway is then derived from the probability versus consequences matrix, modified after CIRIA C552 Table 6.5, given in Table 3.3 below. Note that by definition, no contaminant linkage equates to no risk.

_		Consequence										
	product	Severe	Medium	Mild	Minor							
~	High Likelihood	Very high risk	High risk	Moderate risk	Low risk							
abilit	Likely	High risk	Moderate risk	Low risk	Very low risk							
Probability	Low Likelihood	Moderate risk	Low risk	Low risk	Very low risk							
	Unlikely	Low risk	Very low risk	Very low risk	Very low risk							
	No Linkage	No risk										

This approach assumes an equivalence between probability and consequences and ignores the difficulty that can arise where to probability of occurrence appears to be almost negligible but the consequences are very severe. In such conditions there is a degree of subjectivity in assessing the level of risk and it could be low, moderate or high. Such risks may require specialist consideration beyond the scope of this standard report.

Finally, a description of the classified risks and the likely action required can be determined from Table 3.4 below.

	Description of Classified Risks and Likely Action Required
Very High Risk	A significant contaminant linkage, including actual evidence of significant harm or significant possibility and significant harm, is clearly identifiable at the site (e.g. from visual or documentary evidence) under current conditions, with potential for legal and/or financial consequences for the site owner or other Responsible Person. Remediation advisable based on acute impacts being likely. Immediate action should be considered.
High Risk	A contaminant linkage is identifiable at the site under current and future use conditions. Although likely, there is no obvious actual evidence of significant harm or significant possibility and significant harm under current conditions. Extent of risk is therefore subject to confirmation by investigation and risk assessment and most likely to be deemed significant. Realisation of the risk is likely to present a substantial liability to the site owner or other Responsible Person. Remediation required for redevelopment and may also be required under Part 2A for existing receptors.
Moderate Risk	A contaminant linkage is identifiable at the site under current and future use conditions. However, it is not likely to be a significant linkage under current conditions. It is either relatively unlikely that any such harm would be severe, and if any harm were to occur it is more likely, that the harm would be relatively mild. Actual extent of risk subject to confirmation by additional investigation and risk assessment and most likely to lie between no possibility of harm (under current conditions) and significant possibility of significant harm (under conditions created by new use). Remediation may be required for redevelopment.
Low risk	Potential pathways and receptors exist but history of contaminative use or site conditions indicates that contamination is likely to be of limited extent and below the level of no possibility of harm. It is unlikely that the site owner or other Responsible Person would face substantial liabilities from suck a risk. Precautionary investigations and risk assessment advisable on change of use. Any subsequent remedial works are likely to be relatively limited.
Very Low Risk	No contaminant linkage likely to exist under current or future conditions, but this cannot be completely discounted. If harm is realised, it is likely at worst to be mild or minor. Site not capable of being determined under Part 2A where the local authority inspects the site. No further action recommended.
No Risk	No contaminant linkage exists.

Table 3.4: Description of the Classified Risks and Likely Action Required

3.2 Contaminant Analysis of Samples

The Model Procedures of CLR 11 provide guidance on key information sources with respect to potential contamination arising from past land uses of a site. In particular, the now withdrawn CLR 8 (Environment Agency 2002b), the DoE Industry Profile documents and ISO10381-5 provide good summaries of priority pollutants for UK sites. Additionally, the Environment Agency (2004b) has produced a list of priority pollutants for ecological risk assessment. These documents have been used, with the findings of the Phase 1 investigation, to scope the analyses of chemicals of potential concern. It should be noted that whilst CLR 8 was withdrawn in August 2008 it was not replaced and its findings are still considered useful.

Hydrock considers there to be a minimum requirement for soil chemical analysis, even for greenfield sites, in order to satisfy the 'suitable for use' criterion of the planning regime. This is represented by the 'Hydrock default list of determinands for solids'. The default list is derived from the above guidance, particularly Tables 2.1 and 2.2 of CLR 8, listing potential inorganic and organic contaminants on typical former **industrial** land in the UK.

Since not all redevelopment sites have former industrial land uses, the default list designed to screen for unacceptable risks to property development and future occupiers comprises those substances with human, vegetation and construction materials receptors. The list includes common metals, metalloids and inorganic species, pH, asbestos fibres and screening tests for common organic compound groups which are deemed chemicals of potential concern. Sulfate is a contaminant whose principal receptor is concrete in the ground and is not considered toxic except in extreme conditions. Sulfate analysis is included in the list of geotechnical tests. Some common determinands such as elemental sulfur and sulfide are not included because there is insufficient information available to calculate meaningful assessment criteria.

The Hydrock default list of determinands for water or soil leaching samples is based on the prevailing UK drinking water standards and the environmental quality standards (EQS) values specified by DEFRA (2009) under the UK's obligations under the European Water Framework Directive (WFD). It includes the most common contaminants for use as a screening exercise but does not represent a complete list.

The two Hydrock default lists of determinands are used as a minimum requirement whatever the findings of the Phase 1 investigation. Added to this may be other suites of determinands based on the findings and review of the aforementioned documents.

Assessment is made of all chemicals of potential concern recorded on the site above the laboratory reporting limit. The reporting limits are less than the generic assessment criteria where this is possible. There are two main reasons why this may not be the case.

Firstly, low-level detection may be available using a more detailed analysis method, but this would be disproportionally expensive for routine screening purposes. More detailed testing may be recommended in some instances as an additional phase of investigation once the results of the screening exercise are known.

Secondly, there may be no suitable laboratory method available. In which case it is impossible to give a definitive opinion.

3.3 Generic Risk Assessment Criteria for Human Health

3.3.1 Policy

Generic assessment criteria (GAC) are criteria derived using largely generic assumptions about the characteristics and behaviour of sources, pathways and receptors. These

assumptions will be conservative in a defined range of conditions. The Contaminated Land Exposure Assessment (CLEA) framework uses Soil Guideline Values (SGV) in assessing risks to human health from exposure to soils contaminated with selected contaminants. It has been assumed in this report that the exposure conditions are within the generic conditions used to derive the SGVs.

It should be noted that exceedance of GACs does not automatically mean that the soil is "contaminated". The derivation of GACs includes a number of precautionary assumptions such that non-exceedance will indicate that risk to human health is acceptable and that the land is suitable for use, with regard to the contaminant in question. SGVs are not binding standards, but may be used to inform judgments about the need for action and the selection of remediation standards or target values for individual sites.

However, the legal test for land contamination under the statutory guidance of Part 2A of the Environment Protection Act 1990 (i.e. "significant harm or significant possibility of significant harm") is **unacceptable** intake or direct bodily contact. DEFRA (September 2005 and July 2008) has made it clear that exceedance of a GAC does not necessarily meet this legal test, i.e. exceedance of a GAC does not necessarily equate to unacceptable risk. Consequently, the GACs must be considered as screening values only. The situation was clarified by DEFRA (July 2008) in its guidance on the legal definition of contaminated land and in 2012 by the publication of revised contaminated land statutory guidance. One of the key policy aspects of this revision is to clarify that GACs are only one tool in the decision-making process and that background concentrations and a number of other relevant factors should also be taken into account. The aim is to prevent over-cautious determination of land as being contaminated.

The Environment Agency (2009a) has stated that the Health Criteria Values (HCV) used to derive GACs represent minimal or tolerable risk for long-term human exposure to chemicals in the soil. "Science alone cannot answer the question of whether or not a given *possibility of significant harm* is *significant*, since what is either *significant* or *unacceptable* is a matter of socio-political judgement, and the law entrusts decisions on this to the enforcing authorities (DEFRA July 2008)."

The Health Protection Agency (2009) also describes how HCVs do not represent unacceptable intake and that unacceptable intake is not a toxicological parameter. It further asserts that "unacceptable intake is a policy decision which can only be taken by the local authority." Pointers provided to local authorities in this regard are provided by the following: "The HCVs, and GACs based upon them represent trigger values above which there might be a possibility of significant harm. Whether there is a significant possibility will be linked to factors such as the margin of exceedance, the duration and frequency of exposure, and other site-specific factors."

The 2012 National Planning Policy Framework states that the standard of remediation to be achieved through the grant of planning permission for new development, including permission for land remediation activities, is the removal of unacceptable risk and making sure the site is suitable for its new use. As a minimum, after carrying out the development

and commencement of its use, the land should not be capable of being determined as contaminated under Part 2A. The requirements for planning are, therefore, the same as for Part 2A.

The 2012 contaminated land statutory guidance says that GAC represent cautious estimates of levels of contaminants in soil at which there is considered to be no risk to health or, at most, a minimal risk to health. They may be used to indicate when land is very unlikely to pose a significant possibility of significant harm to human health. They should not:

- be used as direct indicators of whether a significant possibility of significant harm to human health may exist. Also, the local authority should not view the degree by which GACs are exceeded (in itself) as being particularly relevant to this consideration, given that the degree of risk posed by land would normally depend on many factors other than simply the amount of contaminants in soil.
- be seen as screening levels which describe the boundary between Categories 3 and 4 (see below);
- be viewed as indicators of levels of contamination above which detailed risk assessment would automatically be required under Part 2A or, under the planning system, in relation to ensuring that land affected by contamination does not meet the Part 2A definition of contaminated land after it has been developed; nor
- be used as generic remediation targets under the Part 2A regime.

Where it is judged that significant uncertainties remain following assessment against generic criteria, there are two options for the developer: either the implementation of an agreed remedial strategy, or to undertake additional testing and/or a detailed quantifiable risk assessment to determine whether remediation is indeed necessary.

3.3.2 Methodology

The sample analyses are divided into representative data sets for the assessment, based on the conceptual model and taking into account such characteristics as variation in soil properties or historical, existing or proposed land uses. The 'averaging area' is the area of soil to which a receptor is exposed or which otherwise contributes to the creation of hazardous conditions.

The determination of averaging areas is clarified in the CLEA Frequently Asked Questions (30 January 2006) document available from the Agency CLEA web pages. In applying statistical tests, the risk assessor is asking the question "are mean (95 percentile upper confidence limit) soil concentrations within the averaging area equal to, or greater than, the SGV/GAC?" If a garden lies within a larger averaging area, but that averaging area is representative of conditions within the garden, then this is the average concentration a receptor using the garden will be exposed to. An averaging area can, therefore, be larger than a single garden and part of a larger zoned area if:

- contaminant concentrations are within the same statistical population, the sample data being representative of the averaging area and the mean concentration of the averaging area;
- hot spots are treated as separate zones or averaging areas; and
- the sampling strategy takes into account uncertainty (spatial heterogeneity) in contaminant concentration.

The approach taken in this report is to characterize the materials that are likely to form the ground cover in garden areas by zoning the site. Each averaging area has been chosen to describe the area(s) of the site, zoned according to material type and existing conditions, within which assessment against GACs has taken place. As pointed out in P5-066/TR (Environment Agency 2000) and by Nathanail (2004), this is a logical way of investigating a large plot of land that is intended for residential use, particularly if the development layout may not have been finalised.

The original Soil Guideline Values were all withdrawn in August 2008 and the Agency started a programme of publishing replacements using its 'new approach', which involves a number of changes to the way exposure is assessed. This was started using the CLEA 1.04 software. The current version is CLEA 1.06. This programme was put in abeyance when DEFRA started to re-draft the Part 2A statutory guidance.

A significant change in the new approach is to publish SGVs only at 6% soil organic matter (SOM) content. This appears to be counter productive because in cases where the SGV varies with the SOM, the published SGV report states that "at a lower SOM, they may not be sufficiently protective." The SGV introduction report *Using Soil Guideline Values* states that in such circumstances a new GAC can be produced by adjusting the SGV.

Furthermore, the SGVs are no longer published for the residential without plant uptake land use.

Consequently, the approach taken by Hydrock is to take the various input parameters from the SGV reports and produce GACs for various SOM and for the residential without plant uptake land use, using CLEA 1.06. The GACs adopted by Hydrock for the standard CLEA land uses are given in Table 3.5 together with the source of the GAC. The table also lists GACs for open space (see below).

The absence of published SGVs for certain chemicals of potential concern has been addressed by the derivation of GAC using generic assumptions about the characteristics and behaviour of sources, pathways and receptors and the CLEA 1.06 software. Input data have been derived either from published GAC lists (EIC/AGS/CL:AIRE and LQM/CIEH) or by inhouse research of the recommended data sources.

Please note also that CLEA 1.06 allows for other variations, most notably of soil type (9 options) and building type (5 residential options). The defaults are a sandy loam soil, a small

terraced house in the residential setting and a pre-1970s office block in the commercial setting. These are generally conservative and the resultant SGV/GAC are protective of other combinations (unlike the default SOM mentioned above). It is not practical to include all permutations in Table 3.5 and in the cases where specific GACs have been derived, this is referred to in the text of the report and the relevant values included in the assessment tables.

Lead is a special case as the former SGV was not based on the CLEA model, but equations utilising blood lead concentrations. There is currently no guidance on how to risk assess lead and the work-in-progress by the Environment Agency under to derive a new methodology using CLEA is in abeyance. Consequently, the former SGVs for lead have been retained by Hydrock until this has been clarified.

Further details including data sources can be obtained on request. It is Hydrock's policy to continually review GACs and updates are made in response to the latest Government guidance or as more data on the substances becomes available. The date of the last update of the table is indicated.

Table 3.5: Soil GACs Adopted by Hydrock (mg/kg) - on following pages



Updated 20/07/11	Human Health Generic Assessment Criteria (mg/kg)															
Contaminant	Source of GAC	Human health - residential without plant uptake (1%SOM)	Human health - residential without plant uptake (2.5%SOM)	Human health - residential without plant uptake (6%SOM)	Human health - residential with plant uptake (1%SOM)	Human health - residential with plant uptake (2.5%SOM)	Human health - residential with plant uptake (6%SOM)	Human health - allotments (1%SOM)	Human health - allotments (2.5%SOM)	Human health - allotments (6%SOM)	Human health - commercial (1%SOM)	Human health - commercial (2.5%SOM)	Human health - commercial (6%SOM)	Human health - open space (1%SOM)	Human health - open space (2.5%SOM)	Human health - open space (6%SOM)
Under de Defende Onder							Default for SGV			Default for SGV			Default for SGV			
Hydrock Default Suite Arsenic	SGV report + CLEA 1.06	35	35	35	32	32	32	43	43	43	640	640	640	590	590	590
Beryllium	LQM/CIEH + CLEA 1.06	51	51	51	51	51	51	55	55	55	420	420	420	8600	8600	8600
Boron	LQM/CIEH + CLEA 1.06	1000			290			45	45	45	190000			45000		45000
Cadmium Chromium (III)	SGV report + CLEA 1.06 LQM/CIEH + CLEA 1.06	85 630	85 630	85 630	11 630			1.9	1.9 15000	15000	230			860 660000		860 660000
Chromium (VI)	LQM/CIEH + CLEA 1.06	4.3	4.3		4.3		4.3	2.2	2.2	2.2	35	35		4400		4400
Copper	LQM/CIEH + CLEA 1.06	6200	6200	6200	2300		2300	520	520	520	72000			400000		400000
Lead	SGV 10 (old method)	450			450			450	450		750			450		450
Lead	CLEA 1.06 (not yet in use)	220 240		220 240	220 170			590 80	590 80	590 80	5100 3600	5100 3600		11000		11000 9900
Mercury, inorganic Nickel	SGV report + CLEA 1.06 SGV report + CLEA 1.06	130			170			230				1800		38000		38000
Selenium	SGV report + CLEA 1.06	600	600	600	350	350	350	120		120	13000	13000	13000	29000		29000
Vanadium	LQM/CIEH + CLEA 1.06	190	190	190	74	74	74	18	18	18	3200	3200	3200	13000	13000	13000
Zinc Cuantida (franc)	LQM/CIEH + CLEA 1.06	40000 760	40000	40000	3700 750	3700 750	3700 750	620		620	670000	670000 16000		1000000		1000000
Cyanide (free) Phenol	Hydrock + CLEA 1.06 SGV report + CLEA 1.06	310	760 420	760 520	180			2300	2300	2300 280	16000	3200		21000		21000 3000
Acenaphthene	LQM/CIEH + CLEA 1.06	2000	3100	3900	210	480	1000	34	85	200	85000	98000	100000	39000	39000	39000
Acenaphthylene	LQM/CIEH + CLEA 1.06	2000	3000		170			28	69					39000		39000
Anthracene Benz(a)anthracene	LQM/CIEH + CLEA 1.06	20000	22000 5.2	23000 6.2	2300	4900	9200	380	950 5.5	2200 10	520000 89	540000 95		200000		200000
Benz(a)anthracene Benzo(a)pyrene	LQM/CIEH + CLEA 1.06	3.7	5.2	6.2	0.83			2.5	5.5	2.1				13.0		13.0
Benzo(b)fluoranthene	LQM/CIEH + CLEA 1.06	7.0	7.3		5.6	6.5	7.0	3.5	7.4	13	100	100			92	92
Benzo(ghi)perylene	LQM/CIEH + CLEA 1.06	47	47		44			70				660				590
Benzo(k)fluoranthene	LQM/CIEH + CLEA 1.06 LQM/CIEH + CLEA 1.06	10 8.8	10 9.7	10 10	8.5			6.8 2.6	14 5.8	23	140 140			130 130		130 130
Chrysene Dibenz(ah)anthracene	LOM/CIEH + CLEA 1.06	0.86	0.91		0.76			0.76	1.5			140		12.0		12.0
Fluoranthene	LQM/CIEH + CLEA 1.06	970	990	1000	260			52	130	290	23000			8100		8100
Fluorene	LQM/CIEH + CLEA 1.06	1900	2500	2900	160		780	27	67	160	64000	69000		26000		26000
Indeno(123cd)pyrene Naphthalene	LQM/CIEH + CLEA 1.06 LQM/CIEH + CLEA 1.06	4.2	4.4 3.9		3.2 1.5		4.2	1.8 4.1	3.8 9.9	7.1	60 200	61 480		56 13000	56 13000	56 13000
Phenanthrene	LQM/CIEH + CLEA 1.06	840	930	9.3	92			4.1	9.9	23	2200			8100		8100
Pyrene	LQM/CIEH + CLEA 1.06	2300			560			110			54000			20000		20000
TPH fractions			,				,									
TPH ali EC05-EC06 TPH ali >EC06-EC08	LQM/CIEH + CLEA 1.06	30			30			740	1700 5600	3900	300	560	1200	1000000		1000000
TPH all >EC08-EC08	LQM/CIEH + CLEA 1.06	73		370	73 19	160 46		2300 320	770	13000 1700	140 78	320 190	450	1000000 41000		1000000 41000
TPH ali >EC10-EC12	LQM/CIEH + CLEA 1.06	48	120	280	48		280	2200	4400	7300	48			41000		41000
TPH ali >EC12-EC16	LQM/CIEH + CLEA 1.06	24	59	140	24		140	11000	13000				140	41000		41000
TPH ali >EC16-EC35 TPH ali >EC35-EC44	LQM/CIEH + CLEA 1.06 LQM/CIEH + CLEA 1.06	45000 45000	64000 64000	77000 77000	45000 45000	64000 64000	76000 76000	260000 260000	270000 270000	270000 270000	1000000	1000000		81000 81000		81000 81000
TPH aro EC05-EC07	LQM/CIEH + CLEA 1.06	260	480	980	45000		280	200000	270000	270000	1200	2300	4700	180000		180000
TPH aro >EC07-EC08	LQM/CIEH + CLEA 1.06	610	1300	2700	120		610	22	51	120	870	1900	4400	180000		180000
TPH aro >EC08-EC10	LQM/CIEH + CLEA 1.06	33	81	190	27			8.6		51	610	1500		16000		16000
TPH aro >EC10-EC12	LQM/CIEH + CLEA 1.06	180	420	870	69 140		350 590	13	31 57	74	360 36000	900	38000	16000		16000
TPH aro >EC12-EC16 TPH aro >EC16-EC21	LQM/CIEH + CLEA 1.06	1250	1600 1300		250			23	5/	130 260		37000 28000		16000 12000		16000 12000
TPH aro >EC21-EC35	LQM/CIEH + CLEA 1.06	1300	1300	1300	890	1100	1200	370	820	1600	28000			12000		12000
TPH aro >EC35-EC44	LQM/CIEH + CLEA 1.06	1300	1300	1300	890		1200	370	820	1600	28000	28000		12000		12000
TPH >EC44-EC70 VOCs - BTEX & MTBE	LQM/CIEH + CLEA 1.06	1300	1300	1300	1200	1300	1300	1200	2100	3000	28000	28000	28000	12000	12000	12000
Benzene	SGV report + CLEA 1.06	0.27	0.49	1.0	0.08	0.16	0.33	0.017	0.035	0.075	28	50	95	240	240	240
Toluene	SGV report + CLEA 1.06	610	1300		120			22	51	120	870	1900		180000		180000
Ethylbenzene	SGV report + CLEA 1.06	170	380 140	840 320	65	150	350	16	39 67	91 160	520	1200 1100	2800	81000		81000 150000
Xylene, o- Xylene, m-	SGV report + CLEA 1.06 SGV report + CLEA 1.06	60 55			45			28				1100 1500		150000		150000 150000
Xylene, p- (use this for combined m & p)	SGV report + CLEA 1.06	53		290	44			29	74			1400	3200	150000		150000
MTBE	EIC/AGS/CL:AIRE + CLEA 1.06	73		220	49			23			7900		24000			240000
VOCs - other benzenes							}									
Iso-propylbenzene Propylbenzene	EIC/AGS/CL:AIRE + CLEA 1.06 EIC/AGS/CL:AIRE + CLEA 1.06	12 40			11 34		64 190	32 34			390 400	950 980		81000 81000		81000 81000
1.2.4-Trimethylbenzene	EIC/AGS/CL:AIRE + CLEA 1.06	40 0.41			0.35			0.38	0.93		400			81000		
VOCs - chlorobenzenes																
Bromobenzene	EIC/AGS/CL:AIRE + CLEA 1.06	0.91			0.88			3.2			97					19000
Chlorobenzene 1,2-Dichlorobenzene	LQM/CIEH + CLEA 1.06	0.33	0.74	1.7	0.33	0.73	1.7	5.9	14 230	32 540	59	130		59000 1000000	59000 1000000	59000 1000000
1,3-Dichlorobenzene	LQM/CIEH + CLEA 1.06	0.31	0.74	94	0.29	0.70	1.7	0.25	0.61		32				1300	1300
1,4-Dichlorobenzene	LQM/CIEH + CLEA 1.06	42	100	230	30		170	15	37	88	230	540	1300	57000	57000	57000
Hexachlorobenzene	LQM/CIEH + CLEA 1.06	0.20	0.50	1.7	0.20	0.50	1.4	0.18	0.42	0.92	0.2	53		24		24
Pentachlorobenzene 1.2.3-trichlorobenzene	LQM/CIEH + CLEA 1.06 LQM/CIEH + CLEA 1.06	14	21 2.6		5.2		17 6.1	1.2 4.7	3.1 12		43 110	110 270				370 6200
1,2,4-trichlorobenzene	LQM/CIEH + CLEA 1.06	1.1	4.5	<u>0.2</u> 11	1.0		11	4.7	75	180	230			44000		44000
1,3,5-trichlorobenzene	LQM/CIEH + CLEA 1.06	0.23		1.4	0.23	0.57	1.4	4.7	12	28	24			6100	6100	6100
1,2,3,4-tetrachlorobenzene	LQM/CIEH + CLEA 1.06	17	39	84	12	29	62	4.4	11	26	120	300	730	2700	2700	2700
1,2,3,5-tetrachlorobenzene 1,2,4,5-tetrachlorobenzene	LQM/CIEH + CLEA 1.06 LQM/CIEH + CLEA 1.06	0.53	1.3 1.2	3.0	0.49			0.38	0.94	2.2	39	98 49	10	290 41		290 41
VOCs - chloroalkanes & alkanes	LOW CIEFT CLEAT.00	0.51	ş 1.2	2.4	0.30	0.68	1.4	0.064	0.16	0.37	20	49	9/	41	; 41	41
Bromodichloromethane	EIC/AGS/CL:AIRE + CLEA 1.06	0.019	0.034	0.07	0.016			0.016	0.032		2.1			240		240
Bromoform	EIC/AGS/CL:AIRE + CLEA 1.06	5.2	11	2.3	2.8	6.0	13	0.95	2.1	4.6	760	1500	3100	15000	15000	15000



Bank Bank <t< th=""><th>Updated 20/07/11</th><th></th><th colspan="14">Human Health Generic Assessment Criteria (mg/kg)</th><th></th></t<>	Updated 20/07/11		Human Health Generic Assessment Criteria (mg/kg)														
Scheding Control Contro Control <thcontrol< th=""> <th< th=""><th>Contaminant</th><th>Source of GAC</th><th>residential without plant</th><th>residential without plant uptake</th><th>residential without plant</th><th>residential with plant uptake</th><th>residential with plant uptake</th><th>residential with plant uptake</th><th>allotments</th><th>allotments</th><th>allotments</th><th>commercial</th><th>commercial</th><th>commercial</th><th>open space</th><th>open space</th><th></th></th<></thcontrol<>	Contaminant	Source of GAC	residential without plant	residential without plant uptake	residential without plant	residential with plant uptake	residential with plant uptake	residential with plant uptake	allotments	allotments	allotments	commercial	commercial	commercial	open space	open space	
Subscience End Size Add 1 (a) 10 Sole Sole Control Sole																	1000000
1.100000000000000000000000000000000000																	2000
Chief Schultz Cale Schultz Cale Schultz Cale Schultz Cale Schultz Cale Schultz	1.1-Dichloroethane																160000
C) C) C) C) C) C) CO CO <thco< th=""> CO CO CO<td></td><td>LQM/CIEH + CLEA 1.06</td><td>0.0065</td><td>0.0093</td><td>0.016</td><td>0.0053</td><td>0.0080</td><td>0.014</td><td>0.0046</td><td>0.0083</td><td>0.016</td><td>0.71</td><td>1.0</td><td>1.8</td><td></td><td>97</td><td>97</td></thco<>		LQM/CIEH + CLEA 1.06	0.0065	0.0093	0.016	0.0053	0.0080	0.014	0.0046	0.0083	0.016	0.71	1.0	1.8		97	97
			0.23	0.41				0.82					46	92			37000
Spectram COUNTS ALL Count ALL <t< td=""><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td>4700</td></t<>																	4700
S.S.S.S.S.S.S.S.S.S.S.S.S.S.S.S.S.S.S.				0.35								22	40				14000
Sharahaman Expansion <					4.5			1.7									10000
11.3 Control 1.5 (1) 1.5 (2) <td></td> <td>410</td>																	410
11.42 : Section Control 0.000 (1) </td <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td>2.1</td> <td>4.8</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td>11000</td>							2.1	4.8									11000
Specific control Specific contro Specific contro <td></td> <td></td> <td></td> <td></td> <td></td> <td>0.89</td> <td>2.1</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td>4700</td>						0.89	2.1										4700
Displace																	4700
11.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1.1																	4200
11.1 Distribution CASSG. 407. CAS 1.0 0.00 1.4 0.00 1.0 1.0 0.00 1.0 0.00 1.0 0.00 <t< td=""><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td>480000</td><td>480000</td></t<>																480000	480000
Dite Algebrache Discription Discription <thdiscription< th=""></thdiscription<>		EIC/AGS/CL:AIRE + CLEA 1.06		1.8	3.9		1.2	2.7						400			3200
Schwarz Schwarz <t< td=""><td>monoromotinano</td><td>LQM/CIEH + CLEA 1.06</td><td>0.92</td><td>1.6</td><td>3.2</td><td>0.75</td><td>1.3</td><td>2.7</td><td>0.36</td><td>0.70</td><td>1.5</td><td>110</td><td>190</td><td>370</td><td>8100</td><td>8100</td><td>8100</td></t<>	monoromotinano	LQM/CIEH + CLEA 1.06	0.92	1.6	3.2	0.75	1.3	2.7	0.36	0.70	1.5	110	190	370	8100	8100	8100
24 Observation Deak (2) Control (2) <thcontrol (2)<="" th=""> <thcontro (2)<="" th=""> <</thcontro></thcontrol>			· - 3														
All-Dimensional ECADE CLAIGE CALE ALLE 21 10 100 10000 1000 1000 1																	1000
Sampland ECASCA MS - 0.24 105 STO STO <td></td> <td>4200</td> <td></td> <td></td> <td>16000</td>														4200			16000
Stamplator EXAMPLATION														180000			81000
Absticked of the second seco		EIC/AGS/CL:AIRE + CLEA 1.06				81			12	28					81000	81000	81000
3.4.6 Construction LOW Colt LEAK 10.0 D2 110 180 0.87 2.0 4.1 0.31 0.31 0.71 200 4.00 10	4-Methylphenol					81	180	400		28	63						38000
2.4.P Totophene DEAD Col - DLF A 100 DEAD					35		1.3	3.0								320	320
Phase Phase <th< td=""><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td>1000</td></th<>																	1000
Bite Arthorny Instant ECAGCALABE + CLA 16 270 280 280 280 1100 44 120 100 8000 8000 8000 8000 8000 4000		LQM/CIEH + CLEA 1.06	82	110	130	1.4	3.3	7.5	0.22	0.51	1.2	3900	4200	4400	1000	1000}	1000
Bull heary phrade ECASSCLARE < CEATO 4400 4400 4400 4400 4400 4400 440		FIC/AGS/CL:AIRE + CLEA 1.06	2700	2800	2800	280	610	1100	48	120	280	85000	86000	86000	36000	36000	36000
Dr-back probate ELASSCLARF + CLA 1:0 47 11 450 13 67 2.0 5.0 12 47 500 500 6100	Butyl benzyl phthalate																410000
Dis-dock planning ElCASSCL ARF + CLE A 10 3400 3400 3200 3700	Diethyl Phthalate		14	29	65	120	260	570	19	41		14		65	160000	160000	160000
Particisés U U U U			4.7	11													6100
Addres LOWELH + CLEA 168 2.1 2.1 2.2 1.7 2.0 2.1 1.3 2.6 4.0 6.4 6.4 6.4 6.4 6.3 2.4 2.4 2.4 2.3 2.3 2.4		EIC/AGS/CL:AIRE + CLEA 1.06	3400	3400	3400	2300	2800	3100	940	2100	3900	89000	89000	89000	37000	37000	37000
Attrache CondClet + CLEA 1.00 37 32 32 32 0.24 0.68 120 0.70 130 1400 1400 1400 1400 1400 1400 1400 1400 1400 1400 1400 1400 1400 1400 1400 1400 1400 1400 1400 1400 <th< td=""><td></td><td>LOM/CIEH + CLEA 1.06</td><td>21</td><td>21</td><td>22</td><td>17</td><td>2.0</td><td>21</td><td>13</td><td>2.6</td><td>4.0</td><td>54</td><td>54</td><td>54</td><td>23</td><td>23</td><td>23</td></th<>		LOM/CIEH + CLEA 1.06	21	21	22	17	2.0	21	13	2.6	4.0	54	54	54	23	23	23
DDE Hydox + CLA 1.0 1300					32				0.037	0.085						370	370
Dicklosm		Hydrock + CLEA 1.06	1300	1300	1300	860	1100	1200			1300	22000	22000	22000	33000	33000	33000
Decksin LOMCEH + CLEA 106 3.5 3.8 3.9 0.60 1.4 2.2 0.13 0.22 0.73 0.0 91 92 93 93 Encountim - spin LOMCEH + CLEA 106 447 71 72 <th7< td=""><td>DDT</td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td>20000</td></th7<>	DDT																20000
Endocularia-apha LOMC(EH + CLEA 1.06 44 76 110 2.9 7.0 16 0.47 1.2 2.7 2.00 30.00 34.00 160				32				1.3									400
Endosufici-bela LOMOIEH - CLEA 106 53 69 120 28 6.6 15 0.44 1.1 2.6 2000 3200 1800 1600 1600 160 Heachhorsychesanes - brain (nc. Lindarn) LOMOIEH + CLEA 1.06 60 550 650 14 30 0.64 1.5 1100 1100 430 490 44 Heachhorsychesanes - brain (nc. Lindarn) LOMOIEH + CLEA 1.06 60 52 52 1.7 3.9 6.8 0.064 1.5 1100 1100 430 490 44 42 2.2 0.56 550 550 560 550 550 560 550 550 560 550 550 400 2.2 0.26 0.009 0.0097 0.0007 0.0008 0.008 0.008 0.008 0.008 0.008 0.008 0.008 0.008 0.008 0.008 0.008 0.008 0.008 0.008 0.008 0.008 0.008 0.008 0.008																	39
Hauschlorog/obhanames - shaf (nr. Lundam) LOMCEH + LEA 1.06 460 550 650 160 17 39 65 0.26 0.54 1500 1500 6500 651 630 650																	1600
Headenboorgebrhannes - berlag (nc. Lindare) LOMCEH + CLEA 1.06 50 52 52 7.7 3.9 8.5 0.26 0.0.4 1.10 1100 1100 4400 440 440 Headenboorgebrhannes - gramm (n. Lindare) Londarity (Lindare) Softward (Lindare) S					650			100									6500
HeaceActoroglobAsames gamma (inc. Undane) LOM/CEP+ CLEA 1.06 16 22 2 0.50 1.4 0.0007 0.		LQM/CIEH + CLEA 1.06	50	52	52	1.7	3.9	8.5		0.64	1.5	1100	1100	1100	490	490	490
Tail down, furns & DL-PCB (arrial dop.) SV report > CLEA 1.06 0.0099 0.009 0.0097 0.0073 0.008 0.024 0.24	Hexachlorocyclohexanes - gamma (inc. Lindane)	LQM/CIEH + CLEA 1.06	18	22	23	0.58	1.4	3.0	0.089	0.22	0.52	530	550	550	240	240	240
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Updated 20/07/11			Human Health Generic Assessment Criteria (mg/kg)													
Contaminant	Source of GAC	Human health - residential without plant uptake (1%SOM)	Human health - residential without plant uptake (2.5%SOM)	Human health - residential without plant uptake (6%SOM)	Human health - residential with plant uptake (1%SOM)	Human health - residential with plant uptake (2.5%SOM)	Human health - residential with plant uptake (6%SOM)	Human health - allotments (1%SOM)	Human health - allotments (2.5%SOM)	Human health - allotments (6%SOM)	Human health - commercial (1%SOM)	Human health - commercial (2.5%SOM)	Human health - commercial (6%SOM)	Human health - open space (1%SOM)	Human health - open space (2.5%SOM)	Human health - open space (6%SOM)
Dimethyl phthalate	Insufficient data (EIC)	-	-	-			-		-	-	-	-	-	-	-	-
Isopropyltoluene	Insufficient data (EIC)	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
1-Methylnaphthalene	Insufficient data (EIC)	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
2-Methylnaphthalene	Insufficient data (EIC)	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
Sulfur (elemental)	Insufficient data (Hydrock)	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
1,3,5-Trimethylbenzene	Insufficient data (EIC)	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
tert butylbenzene	Insufficient data (EIC)	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
NOTES			·				, ,									(
If >1,000,000 is calculated, 1,000,000 is adopted.							1									1
Red text - liquid at ambient temperature, calculate	ed GAC exceeds saturation value ar	nd highlighed in r	ed in CLEA - sa	turation value a	dopted for GAC											1
Orange text - solid at ambient temperature, calcu	lated GAC exceeds saturation value	and highlighted	red in CLEA - m	nanual calculatio	on not possible :	as only one HC	V - saturated va	pour concentral	tion exceed, so	saturation value	e adopted for G	AC.	1			1
Orange text - solid at ambient temperature, calcu Blue text - solid at ambient temperature, calculate	ed GAC exceeds saturation value an	d highlighted rec	I in CLEA - man	ual calculation r	not possible as	only one HCV -	aqueous solubi	ity exceed, so o	original red-high	lighted value ac	lopted for GAC			1		1
Green text - solid at ambient temperature, calcula	ated GAC exceeds saturation value a	and highlighted re	ed in CLEA - ma	anual calculation	undertaken bu	t result is greate	er than original i	ed-highlighted	value, so origina	al red-highlighte	d value adopte	d for GAC.				1

3.3.3 Exceedance of Saturation Limits

In some instances the CLEA 1.06 model produces GACs with a warning that the value exceeds the saturation value, which is either the solubility of the substance in water or the vapour saturation limit. Limited guidance is given in SR4 (Section 4.12) on how to assess the GAC in these circumstances. Precedence is also set in a number of SGV reports, to date those dealing with the BTEX compounds. These two sets of documentation are contradictory. The original issue of SR4 (CLEA 1.04) (Environment Agency 2009b) gives an example of how to carry out a manual calculation using data for ethylbenzene, whereas the BTEX SGV reports (e.g. Environment Agency March 2009) state that the GAC should be limited to the saturation level. The revised version of SR4 (CLEA 1.05/6) (Environment Agency 2009c) retains the example, but the name ethylbenzene has been removed.

There are three options: to adopt the value as calculated, to limit the GAC to the saturation value, or to undertake a manual calculation as per Section 4.12 of SR4. Again, the guidance is confusing. SR3 (Environment Agency 2009b) cautions against adopting the saturation limit, which is the most conservative, saying that it may be over-conservative. However, this is the approach taken in the BTEX SGV reports.

Clearly, the adoption of a GAC under conditions where the saturation level is exceeded is subjective and professional judgement is involved. With this in mind, the protocol adopted by Hydrock is as follows, and has been derived at by considering the possible values from the three methods given above.

- 1. For substances where the GAC is highlighted in amber in CLEA, this is adopted as the GAC. For substances where the GAC is highlighted in red, the following apply.
- 2. For VOCs including BTEX and the volatile TPH Fractions (less than EC10), the saturation value is adopted in line with the latest recommendations in the BTEX SGV reports.
- 3. For substances which are liquid at ambient temperature, the saturation value is adopted.
- 4. For substances which are solid at ambient temperature, the manual calculation is undertaken provided there are both oral and inhalation HCVs. The result is compared with the red-highlighted GAC and the lower of the two adopted as the GAC. If there is only one HCV and the calculation cannot be performed, the red-highlighted value is adopted as the GAC where the saturation limit exceed the aqueous solubility, but the saturation value is adopted where the saturation limit exceed the saturated vapour concentration.
- 5. In some instances the GACs shows a large difference between different SOM where the saturation value has been taken for, say, 1% SOM and the calculated values for 2.5% and 6% SOM. Whilst this may appear inconsistent on first inspection, the results have been adopted as they are and the difference must be attributed to the physico-chemical influence of organic matter in the soil as modelled by CLEA.

3.3.4 GACs for Public Open Space

One of the main reasons why the originally intended SGVs for public open space have never been developed is the difficulty in defining generic exposure scenarios for people using such land. Consider the difference in exposure to soil contaminants by someone playing rugby on a muddy pitch compared with someone walking a dog in an urban park. The CLEA 1.06 model does not contain an open space land use and in order to calculate GACs with the same degree of assurance it would be necessary to fully define the exposure profile for a number of potential receptors, to use the basic equations contained in the model and to determine which of the receptors is the critical one.

This is clearly a significant body of work, probably involving research into the various exposure scenarios and is beyond the scope of this report. Determining a UK generic land use pattern is probably something only a national government can do as it will involve policy as well as technical decisions.

In the absence of GACs for open space it is common to refer to those for the residential without plant uptake land use and to acknowledge that these are conservative because (a) there is no building and (b) the exposure period is likely to be much less than in a residential setting.

In order to investigate the degree of conservatism in the above approach, Hydrock has endeavoured to produce GACs for open space by using the 'site specific' options in the CLEA 1.06 model, using the residential without plant uptake land use as a starting point. (In fact, the same results are obtained if the allotments land use is taken as the starting point.) It is recognised that these may not necessarily be representative of the most critical exposure scenario, but are considered suitable as a first screening assessment in the absence of a UK formalised approach.

Accordingly, the method has been to use CLEA 1.06 (except for lead) starting with the residential without plant uptake land use and deleting the indoor exposure pathways. The default soil-to-skin adhesion factor and soil/dust ingestion rates for lifetime exposure are taken from Table 3 of the cadmium SGV report (Environment Agency June 2009). Finally, the exposure times were modified. There is little published guidance on exposure times. The New Zealanders assume 350 days/yr and a duration of 30 years, and the Australians 2 hr/day, 365 days/year over 70 years for parklands and recreational land use. The Canadians take 24 hrs/day, 365 days/year but also include residential land use with parklands. Hydrock has adopted the Australian model of 2 hours every day (active) for the entire lifetime, as being a very conservative estimate, but one which could be said to include for the uncertainties in how the land is actually used.

In addition to changing the exposure time to 2 hours per day in CLEA 1.06, it is also necessary to adjust the dust ingestion rates. Since people are asleep for 8 hours and awake for 16 hours, it is assumed that the soil and dust ingestion takes place during the waking hours. Consequently, if someone in on site for 2 hours, this is 2/16 of the waking period. So it is logical to pro-rata the soil and dust intake (i.e. 1/8 times the default). The defaults are

0.1g/day in Age Classes 1-12 and 0.5 g/day in Age Classes 13-18 (the latter being based on the cadmium SGV report, Table 3); these are replaced with 0.0125 and 0.00625, respectively. In addition, the outside soil to skin adhesion factor for the Age Classes 13 to 18 have been changed to 0.3, as per the cadmium SGV report, Table 3.

The calculated GACs are presented in Table 3.5 and are based on the a 75 year lifetime (all 18 CLEA age classes) and the CLEA standard sandy loam soil of pH 7 su.

Contamination by lead is a special case and CLEA is not applicable. The equations in SGV10 (DEFRA and The Environment Agency 2002) have been used instead. There are separate equations for children and adults. The child equation is based on uptake not intake and the GAC is dependent on the slope of the blood lead level versus soil lead concentration graph so there is no scope for adjusting the GAC for different exposure conditions as these are not inputs to the model. The equation for adults does include an exposure factor which is 8.5 hrs/day for 230 days/yr under the standard commercial land use scenario (for 43 years of working life). The exposure factor can be modified, therefore, to represent different exposures (2 hrs/day for 365 days/yr is equivalent to 8.5 hrs/day for 85.9 days/yr and so 85.9 days was used in the equation). The adult open space GAC for 57 years of exposure (75 less 18 pre-adult years) is calculated as 1135 mg/kg but this is greater than the child SGV of 450 mg/kg so 450 is adopted as the GAC for open space.

3.4 Note on PAHs

A number of authors have used to concept of PAH double ratio plots to investigate the possible source of PAHs in environmental samples.

NAVFAC (Appendix A, April 2003) defines three major source type: petrogenic - generated from organic matter in ancient sediments by geologic conditions; pyrogenic – generated by combustion of organic matter (wood, coal, petroleum, wastes etc.); and biogenic – generated by modern biological processes or by diagenetic processes (e.g. oxidation of organic matter). The following broad trends in the data were recognised:

- a ratio of fluoranthene to pyrene (FI/Py) of <1 is indicative petrogenic sources, and of >1 is indicative of pyrogenic sources; and
- a ratio of anthracene to phenanthrene (An/Ph) of <0.2 is indicative of pyrogenic sources and of >0.2 is indicative of petrogenic sources.

Yunker *et al* (2002) carried out a literature study of published PAH ratios for a number of petroleum sources, combustion sources and environmental sources. They identified the following broad trends in the data:

 a ratio of fluoranthene to fluoranthene plus pyrene (Fl/(Fl+Py)) of <0.4 is indicative of petroleum hydrocarbon sources; of 0.4-0.5 is indicative of liquid fossil fuel combustion products; and of >0.5 is indicative of grass, wood and coal combustion products;

- a ratio of benzo(a)anthracene to benzo(a)anthracene plus chyrsene (BaA/(BaA+Ch)) of <0.2 is indicative of petroleum hydrocarbon sources; of 0.2-0.35 is indicative of either petroleum hydrocarbon sources or combustion and of >0.35 is indicative of combustion products;
- a ratio of anthracene to anthracene plus phenanthrene (An/(An+Ph)) of <0.1 is indicative petroleum hydrocarbon sources, but can be emissions from lignite, diesel or oil combustion, and of >0.1 is indicative of combustion sources, but can be diesel, coal or some crude oil hydrocarbons;
- a ratio of indeno(1,2,3)pyrene to indeno(1,2,3)pyrene plus benzo(ghi)perylene (IP/(IP+Bghi)) of <0.2 is indicative of petroleum hydrocarbon sources; of 0.2-0.5 is indicative of petroleum hydrocarbon combustion; and >0.5 is indicative of grass, wood or coal combustion products.

Note that in these authors' study of these and a number of other ratios they cautioned there are exceptions to these generalisations on account of the variability and complexity of, for example, different crude oil sources.

Costa *et al* (2004) and Costa and Sauer (2005) used plots of fluoranthene to pyrene (Fl/Py) against benzo(a)anthracene to chrysene (BaA/Ch), benzo(a)anthracene to benzo(a)pyrene (BaA/BaP) and chrysene to benzo(a)pyrene (Ch/BaP) to distinguish coal tar and creosote contaminants from combustion products they referred to as urban background. They report distinctive areas on the plots relating to the sites being studied. Litton (2006) has also used these ratios to similar effect on other sites.

ALcontrol Laboratories (2006) also uses plots of fluoranthene to pyrene (FI/Py) against benzo(a)anthracene to chrysene (BaA/Ch). Jones (2008) confirms that the following broad trends are derived from unpublished work at the laboratory:

- a ratio of FI/Py of <0.65 is indicative of used engine oil when the ratio of BaA/Ch is higher (approaching 1.40) or other petroleum products when the ratio of BaA/Ch is lower (above about 0.35);
- a ratio of F/Py of 0.65-1 is indicative of petroleum combustion products; and
- a ratio of FI/Py of <1 is indicative of coal when the ratio of BaA/Ch is higher (approaching 1.40) or other combustion soots when the ratio of BaA/Ch is lower (above about 0.35).

It is evident from the literature that if a cross plot is made of two ratios it is often possible to see a separation in samples from different sources and, together with other supporting

information, gain a better understanding of the likely source of the PAHs. Different ratios may give differing degrees of separation and so trying several plots is often useful.

3.5 Note on Petroleum Hydrocarbons

Petroleum hydrocarbon contamination is complex. The type of crude oil, its distillation, processing and blending, and the subsequent weathering in the environment all result in the development of petroleum residues of extreme chemical complexity (Environment Agency, 2003). The laboratory analysis of petroleum hydrocarbons is highly method dependent. In addition to contaminants such as fuels and lubricating oils, the analyses also pick up a range of other chemicals such as PAHs and phenols, together with naturally occurring substances like humic and fulvic matter in organic soils. For example, TPH determination on dried oak leaves can give a result of 18,000 mg/kg of TPH.

TPH can only be used as a surrogate for estimating the petroleum load of a soil if a spill is well defined but is generally not a sound basis for risk management and regulatory control. International approaches for assessing risks from petroleum hydrocarbons focus on dividing the components into groups and assigning toxicologically potency and fate-transport to each group.

Approaches have been developed internationally, one such proposal is discussed by the Dutch National Institute of Public Health and the Environment (RIVM) (Franken *et al* 1999). The approach is broadly to sub-divide the TPH into fractions based on equivalent carbon length for aliphatic (straight chain) and aromatic (cyclic) compounds. The choice of the fractions is based on work carried out by, amongst others, the Total Petroleum Hydrocarbon Criteria Working Group (TPHCWG). The Working Group is guided by a steering committee consisting of representatives from industry, government and academia, with the remit *to develop scientifically defensible information for establishing soil cleanup levels that are protective of human health at petroleum contaminated sites.*

Generic assessment criteria can be developed for each TPH fraction in the same way as they can be for named substances, providing certain assumptions are made regarding the applicability of the data to all the compounds in each fraction. A significant part of the TPHCWG activity has been in determining fraction boundaries to maximize confidence in the eventual criteria.

A modified TPHCWG approach has been adopted in a framework developed by the Environment Agency (2005) for use within the UK. The 13 original TPHCWG fractions have been adopted, with the addition of >EC35-EC44. An undifferentiated (i.e. without aliphatic – aromatic split) fraction of >EC44-EC70 has also been suggested but the Agency says it will be reviewing the need for this in due course, once research has been carried out into the toxicity of these heavy-end products like resins and asphaltenes.

The UK suggested approach to petroleum hydrocarbon risk assessment is summarised as follows:

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 Measure indicator chemicals and compare with their GAC – these are chemicals which are considered as key risk drivers at petroleum hydrocarbon contaminated sites. The chemicals of potential concern depend on the type of hydrocarbon product, but a (nonexhaustive) list has been suggested by the Environment Agency (2005):

Non-threshold: benzene, benzo(a)pyrene, benzo(a)anthracene, benzo(b)fluoranthene, benzo(k)fluoranthene, chrysene, dibenz(a,h)anthracene, indeno(1,2,3,cd)pyrene. **Threshold:** toluene, ethylbenzene, xylene, naphthalene, fluoranthene, phenanthrene, pyrene.

• Measure TPH fractions and compare with their GAC, based on threshold toxicity only.

Aliphatic fractions: >EC5-EC6, >EC6-EC8, >EC8-EC10, >EC10-EC12, >EC12-EC16, >EC16-EC35, >EC35-EC44.

Aromatic fractions: >EC5-EC7, >EC7-EC8, >EC8-EC10, >EC10-EC12, >EC12-EC16, >EC16-EC21, >EC21-EC35, >EC35-EC44.

- **Undifferentiated**: >EC44-EC77 (subject to review and confirmation by Agency).

Hydrock has adopted the first two points from above approach and has developed generic assessment criteria for the TPH fractions up to EC35. These are used for assessment where an appropriate level of sampling and laboratory analysis has been carried out, but cannot be used where more generalised TPH analysis has been scheduled (such as DRO/GRO only).

There is, however, some uncertainty concerning the validity of the additivity check. The Environment Agency (2002a) stated in the now withdrawn CLR 9, Section 4.4, "that it is not valid to simply calculate the sum of the fractions 'soil concentration divided by SGV', and compare this with 1.", because total intake, not just intake from soil, needs to be included. It is assumed that the 2005 document takes this into account and that it is erring on the side of conservatism. Until this is formally resolved, Hydrock will report the additivity check for information, using the LDEQ groupings, but will caution against its use in setting remedial goals without further study or publication of definitive guidance. It is more realistic to carry out the additivity test on individual samples rather than on US₉₅ values for the whole population, because it is unlikely that the TPH profile of the averaging area will be represented by the US₉₅s of every fraction. More likely, a sample high in one fraction will be low in another, particularly where a mixture of products is present in the ground.

The analysis required for the above methodology, using the aliphatic / aromatic split of TPH fractions, is referred to by Hydrock its "**TPH Level 2 suite**" of determinands. In instances where a full numerical risk assessment is not required, Hydrock carries out a screening analysis known as its "**TPH Level 1 suite**" of determinands. The TPH is divided into fractions, but without the aliphatic / aromatic split. This allows a semi-quantitative risk assessment on the basis of taking a worst case condition. The fraction split with the lowest GAC is deemed to apply to the whole fraction. For example, if the Level 1 analysis indicates the presence of >EC8-EC10, the result is compared to the GACs for the aliphatic >C8-C10 and the aromatic >EC8-EC10 fractions. The worst case would be to assume the whole fraction is aliphatic because this is the lower of the two GACs. This is a conservative approach, and if the test is passed, there is no need to proceed further. However, if the test is failed this does not necessarily indicate unacceptable risks and a more detailed risk assessment is required, with the full TPH Level 2 analysis suite.

3.6 <u>Note on Cyanide</u>

Cyanide toxicity is complicated but it is generally accepted that cyanide species exist in 'free' and 'complex' forms. Free cyanide species are toxic and it is generally agreed that free cyanide provides a more scientifically correct basis for the establishment of generic criteria. This approach has been followed in this report.

Metal-cyanide complexes (complex cyanide) are generally not considered toxic but in certain environmental fate reactions it is possible that dissociation may release toxic free cyanide into the water environment. This might occur where complex cyanides are exposed to direct sunlight and photolysis takes place. Such circumstances are considered very rare.

3.7 Note on Polychlorinated Biphenyls

PCBs fall into two groups, the dioxin-like (DL) and the non-dioxin-like (NDL), by virtue of their toxicity.

The Environment Agency methodology for DL-PCBs is included with dioxins and furans in the published dioxins SGV report (Science Report SC050021 / Dioxins SGV). The basis of this report is that because of the additive nature of these substances it is inappropriate to produce individual SGVs. The approach is to obtain speciated analyses of 12 DL-PCBs and, using an Agency spreadsheet, calculate a Hazard Index for a prescribed mixture of substances. SGVs can only be produced for atmospheric fall-out sites where the proportions of the individual substances are assumed to be uniform across the UK according to a table listed in the document.

For potentially industrially contaminated sites (such as where PCBs have escaped from transformers) only a Hazard Index can be produced. This can be converted into a GAC by calculation, but such a GAC is only applicable to conditions where the mixture of substances is unchanged. In effect, Hazard Indices will be calculated for each soil sample and provided these are all less than unity, the site poses no significant risk.

There is not Agency guidance with respect to NDL-PCBs. Hydrock has produced individual GACs for a number of these. A precautionary approach has been taken, in that the NDL-PCBs are assumed to have additive effects and the same approach is taken as with Hydrock's assessment of contamination by TPH fractions. Namely, each substance is compared with its GAC, but there is an additional stage in which a Hazard Index is also calculated. This is similar to the Agency's approach for DL-PCBs, but the Hazard Index calculation is performed at a different stage in the process.

Currently, these two approaches are separate. That is to say, there is no assumption of additvity of effect between DL- and NDL-PCBs. The logic for this is the fact that these two groups where established in the first place on account of their different effects.

The toxicity of the DL-PCBs is far greater than that of the NDL-PCBs. For example, the residential SGV for the full list of dioxins, furans and DL-PCBs under atmospheric fall-out conditions is 0.0087 mg/kg (NB: using only the 12 DL-PCBs in this list gives a GAC of 0.051 mg/kg), whist the lowest GAC for the NDL-PCBs is 0.32 mg/kg under the same exposure conditions. Analyses for DL-PCBs must be undertaken with very low laboratory reporting limits (typically 1ng/kg).

In real life examples, it is almost certain that both forms of PCBs will be present at a site. This is because the marketed products (known as Aroclors) were mixtures of many PCB congeners and they all appear to contain members from both groups (according to literature researched by Hydrock). Perhaps this is why the Agency has only issued guidance on the DL-PCBs.

Logically, if a site contains any PCBs (for example as a 'total' analysis) it is likely to contain DL-PCBs. In which case, the safe concentrations will be very low and can only be confirmed by re-analysing using low detection methods and following the Agency methodology on a sample-by-sample basis. This is effect means that GACs for NDL-PCBs are redundant. The implications of the Agency methodology have yet to be fully understood by the contaminated land community. For example, it would appear that standard laboratory tests for NDL-PCBs are irrelevant. Furthermore, standard reporting limits are far too high, typically 1ug/kg. The only instance where NDL-PCBs become the risk driver at a PCB contaminated site would be if for some reason the DL-PCBs had preferentially degraded.

The Hydrock methodology for PCB risk assessment is to carry out analyses for the 12 DL-PCBs (commonly referred to as the WHO-12) and the 7 most persistent NDL-PCBs (commonly referred to as the ICES-7) at a detection limit of 1 ng/kg (Table 3.6). This is considered conservative because it covers both groups even though the risk driver is most likely to be the DL-PCB group.

The WHO-12 are assessed using the Environment Agency SGV report methodology, to produce a Hazard Index, and the ICES-7 are compared to Hydrock-derived GACs with additivity check. Note that PCB118 appears in both lists and so in assessed under the Environment Agency methodology as a DL-PCB.

Table 3.6: PCB Suites

WHO-12	ICES-7
(Dioxin-Like)	(Most Persistent)
PCB-77	PCB-28
PCB-81	PCB-52
PCB-126	PCB-101
PCB-169	PCB-118
PCB-105	PCB-138
PCB-114	PCB-153
PCB-118	PCB-180
PCB-123	(Non-dioxin-like apart
PCB-156	from PCB-118)
PCB-157	
PCB-167	
PCB-189	

3.8 Note on the Use of Non-UK Assessment Criteria

In rare instances reference to assessment criteria or other trigger values published by other authoritative bodies (other than those concerned with the UK contaminated land regime) may provide background information on the likely degree of contamination of a substance. Trigger levels indicative of naturally occurring concentrations or risk-based guidance from other countries often help place site analysis results into context. It must be remembered that use of non-UK assessment criteria is not in compliance with the UK contaminated land assessment regime given in the Model Procedures. However, these criteria can be of use as an aid to professional judgement and can help in determining a cost-effective and sustainable remedial strategy for a site, in consultation with the regulatory authorities.

3.9 <u>Site Specific Assessment Criteria for Volatile Substances</u>

The CLEA methodology includes the inhalation of indoor vapours where there are occupied buildings in the standard land use scenarios. For volatile substances such as those listed in Table 3.7 the percentage contribution of the indoor vapour pathway to the average daily exposure (ADE) can be seen to be significant (up to 100%). Consequently, if this pathway can be severed by the installation of a suitably designed and installed organic vapour barrier in the buildings only the remaining CLEA exposure pathways need to be considered for the site. Assessment criteria can be calculated for the remaining exposure pathways.

Site Specific Assessment Criteria (SSAC) have been calculated using CLEA UK using the same input parameters etc. as for the Hydrock GACs but with the indoor vapour pathway turned off in the model. The resulting SSACs can be used to inform on risk from these contaminants in the same way as GACs are used, but apply only if suitable membranes are provided and verified.

Table 3.7: Derivation of Site Specific Assessment Criteria for Volatile Substances for CLEA Standard Land Uses *Excluding the Indoor Vapour Pathway* (mg/kg) – on following page(s).



Updated 21/05/10		Hui	nan Health Ger	neric Assessme	ent Criteria (no	indoor vapour	pathway) (mg/	kg)			
Contaminant	Human health - residential without plant uptake, no indoor vapour (1%SOM)	Human health - residential without plant uptake, no indoor vapour (2.5%SOM)	Human health - residential without plant uptake, no indoor vapour (6%SOM)	Human health - residential with plant uptake, no indoor vapour (1%SOM)	Human health - residential with plant uptake, no indoor vapour (2.5%SOM)	Human health - residential with plant uptake, no indoor vapour (6%SOM)	Human health - commercial, no indoor vapour (1%SOM)	Human health - commercial, no indoor vapour (2.5%SOM)	Human health - commercial, no indoor vapour (6%SOM)	Non- Threshold Substance (Y/N)	Notes
VOCs - chloroalkanes & alkanes				0.44	0.01	0.45	540	500	540	N/s s	
Bromodichloromethane Bromoform	26	26	26	0.11	0.21	0.45	510	520	540	Yes No	
Chloroethane	240000	240000	240000	760	1400	2500	2600	3500	5700	No	
Chloroethene (aka vinyl chloride)	1.2	1.2	1.2	0.0037	0.0066	0.012	26	26		Yes	
Chloromethane	220	220	220	0.44	0.86	1.6	1900	2200		No	
1,1-Dichloroethane	17000	17000 11	17000 11	62	120	230	1800	3000	5600 210	No	
1,2-Dichloroethane 1,1-Dichloroethene	3900	4000	4000	0.031 19	0.055 37	0.11 78	200 2200	210 3900	8000	Yes No	
Cis 1,2 Dichloroethene	470	470	480	1.7			3900	6600	9900	No	
Trans 1,2 Dichloroethene	1500	1500	1500	6.2	12		3900 3400	6200	13000	No	
Dichloromethane	270		270 930	0.68	1.2	<u>2.3</u> 17	7300	9000	9000 11000	No	
1,2-Dichloropropane Hexachloroethane	850 44	44	930	4.1	8.1	8.6	1200 910	2100 920		No No	
Tetrachloroethene	1200	1200	1200	11	24	55	420	950	26000	No	
1,1,1,2-Tetrachloroethane	500	500	500	5.2	12	28	10000	11000	11000	No	
1,1,2,2-Tetrachloroethane	500	500	500	2.7	5.9	13	11000	11000		No	
Tetrachloromethane Trichloroethene	120 450	120 460	130 460	1.1 2.8	2.4 6.3		1500 1500	2500 3200	2500 9000	No Yes	
1,1,1-Trichloroethane	52000	52000	53000	320	700	14	1400	2900	6400	No	
1,1,2 Trichloroethane	350	350	350	1.9	4	8.9	7200			No	
Trichloromethane	610	610	610	2.4	4.7	9.7	5200	9100	20000	No	
Other phenois & chlorophenois 2-Chlorophenoi	1								1	No	Additive, total chlorophenols
2,4-Dichlorophenol		-	-	-		-	-	-	-	NO	Additive, total chlorophenols
2,4-Dimethylphenol	-	-	-	-	-	-	-	-	-	No	
2-Methylphenol	-	-	-	-	-	-	-	-	-	No	Additive, total cresols
3-Methylphenol		-		-			-		-	No	Additive, total cresols
4-Methylphenol Pentachlorophenol	-	-	-	-		-	-	-	-	No No	Additive, total cresols
2,3,4,6-Tetrachlorophenol	-	-	-		-		-	-	-	No	Additive, total chlorophenols
2,4,6-Trichlorophenol	-	-	-	-	-	-	-	-	-	No	Additive, total chlorophenols
Phthalates											
Bis (2-ethylhexyl) phthalate		-	-	-	-	-	-	-	-	No	
Butyl benzyl phthalate Diethyl Phthalate	-	-	-	-	-		-	-	-	No No	
Di-n-butyl phthalate	-	-	-	-	-	-	-	-	-	No	
Di-n-octyl phthalate	-	-	-	-	-	-	-	-	-	No	
Pesticides	1								1		
Aldrin Atrazine	-	-	-	-	-	-	-	-	-	No No	
DDE	-	-	-	-	-	-	-	-	-	No	
DDT	-	-	-	-	-	-	-	-	-	No	
Dichlovos	-	-	-	-	-	-	-	-	-	No	
Dieldrin Endosulfan - alpha			-	-						No No	
Endosulfan - beta	-	-	-	-	-	-	-	-	-	No	
Hexachlorocyclohexanes - alpha (inc. Lindane)	-	-	-	-	-	-	-	-	-	No	
Hexachlorocyclohexanes - beta (inc. Lindane)	-	-	-	-	-	-	-	-	-	No	
Hexachlorocyclohexanes - gamma (inc. Lindane) Dioxins, furans & dioxin-like-PCBs	-	-	-	-	-	-	-	-	-	No	
Total dioxins, furans & dioxin-like-PCBs Non-dioxins, furans & DL-PCB (aerial dep.)	-	-	-	-	-	-	-	-	-	No	
PCB-28	-	-	-	-	-	-	-	-	-	No	
PCB-52	-	-	-	-	-	-	-	-	-	No	
PCB-101 PCB-138	-		-			-	-	-	-	No	
PCB-138 PCB-153	+	-		-				-		No No	
PCB-180	-	-	-	-	-	-	-	-	-	No	
Explosives					I		1	1	I		
HMX RDX		-	-	-	-	-	-	-	-	No	
RDX 2,4,6-Trinitrotoluene		-			-		-	-	-	No Yes & No	
Other metals	1										
Antimony	-	-	-	-	-	-	-	-	-	No	
Barium Marguna alemental	-	-	-	-	-		-	-	-	No	
Mercury, elemental Molybdenum	4.3	- 11	- 26	4.3	- 11	- 26	4.3	- 11	- 26	No No	
Other organics											
Biphenyl		-	-	-	-	-	-	-	-	No	
Carbon disulphide	4300	4300	4400	32	67	150	2100	4200	75000	No	
2,4-Dinitrotoluene 2,6-Dinitrotoluene	<u>+</u>	· · ·						· · ·	-	No No	
Hexachloro-1,3-butadiene	- 8.9	- 8.9	- 8.9	- 1.4	- 2.8	- 4.7	- 290	- 290	- 290	NO	<u> </u>
Mercury, methyl	-	-	-	-	-	-	-	-	-	No	
Styrene	1100	1100	1100	10	24	55	23000	23000	23000	No	
Tributyl tin oxide		-	-	-	-	-	-	-	-	No	
2-Chloronaphthalene		-	-	-	-	-	-	-	-	No	1

3.10 <u>Determination of Contaminated Land Under Part 2A of the Environmental Protection</u> <u>Act 1990</u>

The legal test for land contamination under the statutory guidance of Part 2A of the Environment Protection Act 1990 (i.e. "significant harm or significant possibility of significant harm") is **unacceptable** intake or direct bodily contact.

The situation was clarified by DEFRA (July 2008) in its guidance on the legal definition of contaminated land.

Part 2A does not prescribe number-based thresholds because it would be very difficult to produce numbers which are meaningful and proportionate, given the lack of scientific information about many substances and the site specific nature of risks. Instead, it relies on local authorities to assess risks posed on individual sites, then decide whether (in their view) the risks represent SPOSH, and thus whether land qualifies as *contaminated*.

The intention of the approach is that local authorities can use their judgement to ensure that Part 2A focuses on the SPOSH it was designed to address, whilst avoiding unnecessary burdens on land where contaminants may be present but there is no SPOSH.

In making Part 2A decisions, local authorities are likely to face some difficult decisions caused by uncertainty on the nature of risks. But they should be confident in exercising their judgement on the basis of available information. Part 2A clearly leaves judgements about what constitutes a SPOSH to local authorities, and it is up to them to make decisions.

GACs are not proxy thresholds for SPOSH, and should not be used as such. They describe levels (based on cautious estimates and assumptions in hypothetical example situations) at which concentrations of contaminants in soil may cease to pose *no appreciable/ minimal* risk. They do not seek to describe levels at which there might be a SPOSH.

Thus, if a GAC is exceeded, the assessor will usually need to conduct a detailed quantitative risk assessment to discover whether there is a *possibility of significant harm* and, if so, the nature of that risk. Whether or not SPOSH exists will depend on the results of risk assessment, the existence and nature of any pollutant linkages, and (ultimately) the judgement of the local authority.

As a general guide:

- (i) For substances where there is a GAC, the more the GAC is exceeded, the more likely it is that an authority should consider the risks to be SPOSH.
- (ii) Generally, the cautious nature of GACs means that local authorities may conclude that SPOSH is unlikely to exist at concentrations close to GACs.

- (iii) In some cases, land with concentrations of contaminants which marginally exceed a GAC (say, up to a few times the GAC) might give rise to SPOSH if, for example, the receptor is particularly sensitive; or if further assessment finds that exposure is higher than that estimated in the GAC; or if there is little uncertainty in the underlying toxicology and HCV.
- (iv) In other cases a GAC may be exceeded by tens of times and there might be no SPOSH (e.g. if further assessment found that exposure was much lower than that estimated using the GAC).

In view of the above, Hydrock has not to attempted to derive numerical SPOSH concentrations, but to use GACs as screening values. Where GACs are exceeded, it is recommended that the linkages and the uncertainties in the data are reviewed in consultation the regulatory authority to aid its judgment on determination.

A possible next phase would be to refine the generic risk assessment with a detailed risk assessment. This would involve using site-specific input parameters relevant to the particular site, in the CLEA model.

Revised contaminated land statutory guidance was published by DEFRA in 2012 with respect to Part 2A. The Act itself is unchanged. A new four category test (and associated classifications) has been introduced to ensure a high standard without being excessive. The aims are to make the regime target higher risk sites more efficiently, remove excessive cost burdens and facilitate the development of technical tools to increase consistency over time. This includes supporting non-technical guidance including a possible framework to aid in deciding into which of the proposed four new Categories of land a site should be placed.

Conversely, the regime is not intended to intervene where there is only a low level of risk, particularly in cases where it is difficult to demonstrate anything other than a very small hypothetical risk, as might be the case with vast swathes of land.

DEFRA states that there is a need for a more pragmatic approach. In practice, deciding when regulatory intervention is justified involves making decisions about when to act on a wide spectrum of risk, with varying levels of uncertainty over the precise nature of the risks. A number of the changes are intended to clarify when land is "contaminated land". These are most likely to affect the assessment and remediation of contaminated land and are listed below.

 Statutory explanation of broad objectives of the regime to explain that regulators should seek a reasonable balance between dealing with unacceptable risks whilst ensuring that burdens on businesses and society are manageable and sustainable. The regime should be seen as an option of last resort; that land is in effect "innocent until proven guilty". This should give greater clarity for all concerned on what the regime seeks to achieve, and what it seeks to avoid.

- 2. Local Authorities to produce risk summaries before land may be determined as "contaminated". Summaries must be understandable to non-experts to provide greater transparency and accountability. Easier for all involved to understand what local authority considers risks to be. It should be easier for Local Authorities managers, lawyers and councillors to be involved in decision making, particularly more difficult sites where wider socio-economic effects need to be take into account. Easier to share experience between Local Authorities leading to greater consistency in decision making.
- 3. Clarification of the legal test of significant harm to human health to mean serious unhealthy conditions of the body or part of it, and not minor/trivial complaints. This is unlikely to have a major effect because, to date, no site in England and Wales has been determined on grounds that significant harm to human health has actually been caused. However, greater clarity on the meaning of significant harm is likely to help clarify the related legal test of significant possibility of significant harm.
- 4. Explanation of how to decide when land is (and is not) "contaminated land". A new four category test which recognises the spectrum of risk encountered by assessors, and the reality that some sites are clearly contaminated land (Category 1), some clearly are not (Category 4), and others need more detailed consideration before a decision can be taken (Categories 2 and 3). Greater clarity that decision making is a two stage process in which the regulator must first understand the risk before deciding whether the risk is sufficiently high to justify regulatory intervention. The aim is to create legal certainty around what definitely is, and is not, contaminated land, whilst leaving Local Authorities with discretion to exercise local judgement on less straightforward land.
- 5. Category 4 will include normal background levels of contamination unless there is some exceptional reason to consider otherwise. Clarification that land at SGV/GAC levels is likely to be well into Category 4. Statutory backing for the sector to develop new tests to describe the top of Category 4 (including the production of Category 4 Screening Levels). This should provide clarity on when land will not be caught, reduced uncertainty and costs for landowners and businesses and faster decision making on non-problematic land.
- 6. Clarify the status of GACs and how they should (and should not) be used including a legal backing for the use of robust GACs produced by reputable, non-governmental, organisations within the sector (LQM/CIEH, EIC/AGS/CL:AIRE). Backing the development of new GACs (or similar tools) as might be developed by the sector to help implement the new Guidance. Specific legal backing for the current set of SGVs/GACs, and clarity on how they can (and cannot) be used.
- 7. Category 1 land is clearly caught by the regime when there is clear evidence of an unacceptable risk (e.g. similar land is known to have caused significant harm). This should give clarity on when land is definitely "contaminated land", and help frame the spectrum of risk raised by land contamination.

- 8. New category of land under which Local Authorities would decide whether a site is in Category 2 (contaminated land) or Category 3 (not contaminated land). The new test would rest on whether or not the local authority believes there is a strong case for regulatory action, taking account of the scientific evidence, the objectives of the regime, and other factors. The local authority would start by considering health risks alone, and if they clearly tend towards the Category 4 or the Category 1 the decision could be taken at this point. However, if this does not lead to a decision, the local authority would consider wider socio-economic factors (e.g. cost, views of local people, etc) before deciding. If the local authority still cannot decide, the default decision is that the site is not contaminated land.
- 9. Reduce "regulatory creep" (excessive remediation of land forced by regulatory uncertainty) with greater clarity on what the enforcing authority can "reasonably" require by way of remediation. Clarity that SGVs/GACs must not be used as "one size fits all" remediation requirements; and that Part 2A can only be used to force remediation to a level where land is no longer contaminated land (i.e. to a point where land is in Category 3), but it should not be used to force remediation beyond this point.
- 10. Guidance on the process of risk assessment: the need to take a strategic approach; the aim of dismissing low risk sites as soon as possible in order to focus on finding higher risk sites; and the general need to ensure that risk assessment is conducted in a timely and efficient manner. Clarify that in considering possible future risks the local authority should consider likely future situations (e.g. rather than hypothetical worst possible case situations). Recognise that in practice there is often a need for authorities to bring in external experts and act in accordance with their advice. Recognise that scientific and technical uncertainty is an inevitable part of contaminated land risk assessment, and set out broadly how regulators should deal with it. It is important that this is recognised in the Guidance to support the regulators who have to make decisions in the face of uncertainty.

In deciding whether or not a significant possibility of significant harm to human health exists, the local authority should first understand the possibility of significant harm from the relevant contaminant linkage(s) and the levels of uncertainty attached to that understanding, before it goes on to decide whether or not the possibility of significant harm is significant.

The term "possibility of significant harm" means the risk posed by one or more relevant contaminant linkage(s) relating to the land. It comprises:

- the estimated likelihood that significant harm might occur to an identified receptor, taking account of the current use of the land in question; and
- the estimated impact if the significant harm did occur i.e. the nature of the harm, the seriousness of the harm to any person who might suffer it, and (where relevant) the extent of the harm in terms of how many people might suffer it.

Having completed its estimation of the possibility of significant harm, the local authority should produce a risk summary.

The decision on whether the possibility of significant harm being caused is significant (SPOSH) is a regulatory decision to be taken by the relevant local authority. In deciding whether the possibility of significant harm being caused is significant, the authority is deciding whether the possibility of significant harm posed by contamination in, on or under the land is sufficiently high that regulatory action should be taken to reduce it, with all that would entail.

In deciding whether or not land is contaminated land on grounds of significant possibility of significant harm to human health, the local authority should use the four categorisations.

The decision between Categories 2 and 3 is a positive legal test, which means that the starting assumption should be that land does not pose a significant possibility of significant harm unless there is reason to consider otherwise. Category 3 may include land where the risks are not low, but nonetheless the authority considers that regulatory intervention under Part 2A is not warranted.

The local authority should first consider its assessment of the possibility of significant harm to human health, including the estimated likelihood of such harm, the estimated impact if it did occur, the timescale over which it might occur, and the levels of certainty attached to these estimates. If the authority considers, on the basis of this consideration alone, that the strong case does or does not exist, the authority should make its decision on whether the land falls into Category 2 or Category 3 on this basis regardless of any other factors.

However, if the authority considers that it cannot make a decision, it should consider other factors which it considers are relevant, including:

- The likely direct and indirect health benefits and impacts of regulatory intervention including benefits of reducing or removing the risk posed by contamination, any risks from contaminants being mobilised during remediation and any indirect impacts such as stress-related health effects that may be experienced by affected people, particularly local residents. If it is not clear to the authority that the health benefits of remediation would outweigh the health impacts, the authority should presume the land falls into Category 3 unless there is strong reason to consider otherwise.
- The authority's initial estimate of what remediation would involve; how long it would take; what benefit it would be likely to bring; whether the benefits would outweigh the financial and economic costs; and any impacts on local society or the environment from taking action that the authority considers to be relevant.

Deregulatory change to definition of contaminated land as it relates to water pollution.

DEFRA will commence Section 86 of the Water Act 2003 so that in future this would only be the case if there is significant pollution of controlled waters or significant possibility of such

pollution. To explain how to decide whether or not "significant" pollution is being caused, the Statutory Guidance introduced new Category 1-4 tests similar to those for deciding when there is a significant risk to human health as described above. There will be new technical guidance produced by the Environment Agency. In practice, this change is likely to have little effect on the practical implementation of the Part 2A regime because the Environment Agency has already been prioritising sites likely to meet the new "significance" test.

The 'pollution of Controlled Waters' means the entry into controlled Waters of any poisonous, noxious or polluting matter or any solid waste matter. Given that the Part 2A regime seeks to identify and deal with significant pollution (rather than lesser levels of pollution), the local authority should seek to focus on pollution which: (i) may be harmful to human health or the quality of aquatic ecosystems or terrestrial ecosystems directly depending on aquatic ecosystems; (ii) which may result in damage to material property; or (iii) which may impair or interfere with amenities and other legitimate uses of the environment.

In deciding whether significant pollution of Controlled Waters is being caused, the local authority should consider that this test is only met where it is satisfied that the substances in question are continuing to enter controlled Waters; or that they have already entered the waters and are likely to do so again in such a manner that past and likely future entry in effect constitutes on-going pollution.

3.11 Generic Risk Assessment Criteria for Risk to Plants

Soil contaminants, if present at sufficient concentrations, can have an adverse effect on the plant population. Phytotoxic effects can be manifested by a variety of responses, such as growth inhibition, interference with plant processes, contaminant-induced nutrient deficiencies and chlorosis (yellowing of leaves). All chemicals are probably capable of causing phytotoxic effects. Thus the phytotoxic potential of substances is dependent on the concentrations capable of having adverse effects on plants and the concentrations likely to be found at contaminated sites. Phytotoxicity is a difficult parameter to quantify given that experimental techniques vary widely and variations exist in plant tolerances, soil effects and synergistic/antagonistic reactions between chemicals.

Contaminants may be taken up and accumulated by plants through a range of mechanisms. The principal pathways are active and/or passive uptake through the plant root, adsorption to root surfaces and volatilisation from the soil surface followed by foliar uptake. After plant uptake, contaminants may be metabolised or excreted, or they may be bioaccumulated.

Many of the substances capable of adversely affecting vegetation exert this effect because of their water solubility, a characteristic that could result in their transport from contaminated sites into adjacent locations where the chemical may generate a phytotoxic response. This could be important if, for example, the adjacent site has important conservation status.

Whilst many contaminants may be phytotoxic, data are limited. Some heavy metals are essential as trace elements for plant growth but may become toxic at higher concentrations.

Toxicity may be displayed in many forms, including signs of stress such as reduction in growth or yellowing of the tissue. The concentration in soil at which substances become phytotoxic depend on a range of factors including plant type, soil type, pH, the form and availability of the contaminant and other vegetation stress factors that may be present (such as drought).

Hydrock has carried out a review of a number of current and former guidance documents and other texts on phytotoxicity. It is not possible to produce a definitive list of phytotoxic substances on account of the variables mentioned above. However, a number of metals are repeatedly cited as commonly occurring priority pollutants. As a result, the following list is adopted as Hydrock's indicators of the potential for phytotoxicity: As, B, Cr, Cu, Ni and Zn.

As the CLEA framework is a risk based approach, applied to humans, an alternative strategy is required to assess the risk to plants from substances that are phytotoxic. Reference to published criteria and background concentrations can help put site data into context.

Published assessment criteria for the protection of plant life from a number of countries are given in Table 3.8. Also included in the table are some measures of natural background concentrations in typical soils.

The most authoritative source is the British Standard for topsoil, but this only lists three elements. CLR 11 states that the ICRCL Guidance Note 70/90 can be used for initial screening criteria. This approach has been adopted by Hydrock where BS3882 is lacking, but where an ICRCL 70/90 criterion is lacking, the lowest criterion in Table 3.8 from, firstly MAFF, and, secondly, another country has been adopted. The adopted criteria are highlighted in Table 3.8. The MAFF value of 250 mg/kg has been chosen for As over the ICRCL value of 50 mg/kg as MAFF explains the 50 is applicable to vegetables and human health, whereas 250 is applicable to the plants themselves.

Reference	As	В	Cr (total)	Cr (III)	Cr (VI)	Cu	Ni	Zn
Published Assessment (Published Assessment Criteria (mg/kg)							
						200 (pH>7)	110 (pH>7)	300 (pH>7)
British Standard for topsoil (BS3882:2007)						135 (pH 6-7)	75 (pH 6-7)	200 (pH 6-7)
						100 (pH 5.5- 6.0)	60 (pH 5.5- 6.0)	200 (pH 5.5- 6.0)

Table 3.8: Published Assessment Criteria and Natural Background Concentrations for Phytotoxic
Elements (mg/kg)

Reference	As	В	Cr (total)	Cr (III)	Cr (VI)	Cu	Ni	Zn
MAFF Code of Good Agricultural Practice for the Protection of Soil (1998)	250		(1014)	unlikely to be toxic except in v low pH. 400 for sites containi ng sewage sludge		500 (grass) but may fall to 250 for clover and sensitive species (at pH≥6)	110 (pH>7) 75 (pH 6-7) 60 (pH 5.5- 6.0)	1000 (clover & grass at pH 6), may fall to 300 for sensitive species (at pH 6- 7)
Australian Guideline B(1) (1999), Interim Urban Ecological Investigation Level (EIL). Soils not generally considered phytotoxic below these EILs.	20			400	1	100	60	200
Considered toxic to plants - Ponnamperuma <i>et al</i> (1979)		5 (hot water soluble)						
Dutch ecotoxicological intervention value (Swartjes 1993 & 1994) *	40	7	230			190		
Alberta Environment (1990) Tier 1 (draft) *	10 acid sandy soils			600 acid sandy soils	25 acid sandy soils	130 acid sandy soils		
Ontario MoE (1989) *	20 acid sandy soils 25 clay soils							
ICRCL 59/83 (1987) now withdrawn for human health assessment		3 (hot water soluble)				130	70	300
ICRCL 70/90 (1990) threshold trigger value	50				25	250		1000
New Zealand guidelines for timber treatment sites (1997), estimated based on Cu bioavailability *						500- 1000 clay soils		
New Zealand guidelines for timber treatment sites (1997), soil criteria for protection of plant life (residential/agricultural setting)	10-20	3 (soluble)		600	25	130		
Natural Background Con	centration	s (mg/kg)						
Dutch background level (target value) (VROM 2000)	29		100			36	35	140
UK ICRCL 42/80 (2nd ed. 1983) - Normal conc. In agricultural soil	0.1-40	2-100	5-500			2-100	5-500	10-300

Reference	As	В	Cr (total)	Cr (III)	Cr (VI)	Cu	Ni	Zn
UK ICRCL 70/90 (1st ed. 1990) - Typical range (and mean) in agricultural soils	2.3 - 53 (11.0)					5.8-62 (19) [1.2-19 4.9) extracta ble]		29-210 (78.1) [1.5-21 (5.6) extracta ble]
Canadian assessment criteria (i.e. background) (CCME 1991)	5	1(hot water soluble)	20		2.5	30		60
New Zealand timber sites (1997) – background	2-30							
Australian Guideline B(1) (1999), typical background levels	1-50		5-1000			2-100	5-500	10-300
* cited in New Zealand Ministry for the Environment (1997) timber treatment chemicals guidelines.								

3.12 Generic Risk Assessment Criteria for Controlled Waters

The following aquifer definitions are adopted.

- **Principal aquifers** These are layers of rock or drift deposits that have high intergranular and/or fracture permeability meaning they usually provide a high level of water storage. They may support water supply and/or river base flow on a strategic scale. In most cases, principal aquifers are aquifers previously designated as major aquifer.
- Secondary aquifers These include a wide range of rock layers or drift deposits with an equally wide range of water permeability and storage. Secondary aquifers are subdivided into two types:
- Secondary A permeable layers capable of supporting water supplies at a local rather than strategic scale, and in some cases forming an important source of base flow to rivers. These are generally aquifers formerly classified as minor aquifers; and
- Secondary B predominantly lower permeability layers which may store and yield limited amounts of groundwater due to localised features such as fissures, thin permeable horizons and weathering. These are generally the water-bearing parts of the former non-aquifers.
- Secondary undifferentiated has been assigned in cases where it has not been possible to attribute either category A or B to a rock type. In most cases, this means that the layer in question has previously been designated as both minor and non-aquifer in different locations due to the variable characteristics of the rock type
- **Unproductive strata** These are rock layers or drift deposits with low permeability that have negligible significance for water supply or river base flow.

The European Water Framework Directive (2000/60/EC) (WFD) and its daughter Directives establish a consolidated way of controlling water quality. The UK Government has set out a timetable for the adoption of the WFD which formalises the way in which the quality of surface water and groundwater are to be assessed. This is set out in *The River Basin Districts Typology, Standards and Groundwater threshold values (Water Framework Directive) (England and Wales) Directions 2009.* The regime was established in December 2009, to be fully operational by December 2012. This involves the introduction of new regulations and water quality targets (WQT). The Environment Agency (July 2008) has issued a revised Groundwater Protection Policy (known as GP3). Parts 1-3 contain the high level policy, the technical background and the tools to be used. Part 4 sets out the legal framework and the detailed policies. There are a number of support documents produced by the UK Technical Advisory Group on the Water Framework Directive (UKTAG), including Paper 11B(iii) (September 2008) on standards for regulation.

A groundwater body is defined as groundwater in an aquifer capable of supporting an abstraction of 10 m³/day or 50 people over a sustained period under the WFD. Groundwater bodies are a strategic resource, even if there is no current abstraction. Lesser amounts of groundwater in an aquifer are not considered as receptors in their own right, but may still be pathways to other receptors such as surface water bodies or aquatic ecosystems.

Hydrock has followed the guidance in these documents as far as possible, but because the regime is being introduced over a period of several years, it will not always be possible to 'second guess' the exact requirements of the Agency during the transitional period and so early liaison is recommended.

One of the main objectives of the Agency is to 'prevent or limit' inputs of substances. The List I and List II groupings of substances under the 1980 Groundwater Directive (GD) no longer apply. Substances are instead treated as either 'hazardous substances' (initially broadly equating to the former List I) or non-hazardous pollutants' (analogous to the former List II, but potentially applying to all other pollutants).

The 'limit' objective refers to limiting any inputs of all other pollutants into groundwater so as to prevent pollution, deterioration in status or sustained upward trends. The existing GD already implements the limit objective, but for a limited range of substances (List 2) and activities. The major change in regulation is that now the WFD covers all other pollutants.

For practical purposes, the Agency interprets prevention of inputs of hazardous substances in relation to former List 1 substances and any other substances which meet the criteria for persistence, toxicity and bioaccumulation taking into account those substances listed in WFD Annex VIII. List I substances are effectively a (large) subset of the potentially wider group of hazardous substances.

Note, however, that the 'prevent' objective applies to active inputs such as industrial discharges and *de minimus* concentrations are set as a series of minimum reporting values (MRV). Inputs to controlled waters from contaminated land sites are classed as passive inputs under the WFD and, as such, are regulated under the Agency's 'limit' objective.

Acceptable water quality targets are defined for protection of human health (based on drinking water standards (DWS)) and for protection of aquatic ecosystems (environmental quality standards (EQS)).

There are no longer finite lists of substances over which control may be exercised under the 2006 WFD and the 2009 Groundwater Regulations. All substances which are not determined to be hazardous are potentially non-hazardous pollutants. This enables control to be exercised over polluting substances which have hitherto been beyond control purely because, regardless of their impact, they were not listed in the 1980 Groundwater Directive. In practice the Agency will need to deal with substances which are current priorities of concern. It is clearly not necessary to expand the field to include all other substances in all circumstances unless they are liable to cause pollution.

The final say lies with the Environment Agency and whilst the indicator substances analysed for by Hydrock in this report may be indicative of the likely risk of pollution of Controlled Waters, this report may not be definitive and the Agency may require additional work.

The definition of pollution is *"the direct or indirect introduction, as a result of human activity, of substances or heat into the air water or land which may be harmful to human health or the quality of aquatic ecosystems or terrestrial ecosystems directly depending on aquatic ecosystems, which result in damage to material property, or which impair or interfere with amenities and other legitimate uses of the environment."*

Pollution equates to harm. In order to protect receptors there is a regulatory regime. This involves setting an environmental standard at the receptor (i.e. minimum acceptable water quality). In recognition that pollutants may degrade *en route* to the receptor it is possible to set a limit value at the source of the pollution and compliance values at locations along the pathway, such that water reaching the receptor does not exceed the environmental standard. By definition, the target value is greater than or equal to the compliance value, which in turn is greater than or equal to the environmental standard, depending on the amount of degradation expected. This concept is used in the Remedial Targets Methodology (Environment Agency 2006) to determine how land contamination impacts on groundwater and surface water quality.

The applied environmental standards vary with the hydrogeological conditions and the perceived value of the water resource, and are subject to local assessment by the Agency. Note that protection of Controlled Waters may involve work over and above that required for 'suitable use' of a site for the proposed development.

Note also that Article 6.3(e)(ii) of the WFD enables the regulatory authorities to exempt measures from the prevent and limit requirements where it would be disproportionately costly to remove or control the further movement of pollutants that are already in the ground. Where a continuing source that has given rise in the past to land contamination this must be brought under control to prevent further unacceptable inputs to groundwater, but it is clear that the extent is limited by what is considered to be 'reasonableness'.

This report provides an initial assessment of the risks of pollution of Controlled Waters using water quality targets (WQT) as screening values. These are the drinking water standards (DWS) and the environmental quality standards (EQS), the latter designed to protect the surface water ecosystems. EQS are available for inland surface waters (freshwater) and other surface waters (transitional and marine).

DWS are given in the Water Supply Regulations 2010 (which amends to Water Quality (Water Supply) Regulations 2000, Schedule 1, Table B, Part 1 (Directive requirements) and Part 2 (national requirements)). Where no UK or EU drinking water standard exists, reference is made to the World Health Organization (2011).

The primary list of EQS has been published recently in European Directive 2008/105/EC, Annex I, Part A and these have been adopted by DEFRA in *The River Basin Districts Typology, Standards and Groundwater threshold values (Water Framework Directive) (England and Wales) Directions 2009* (Part 5 Priority Substances).

In addition, each Member State has to define country-specific substances and their EQS. Those adopted by the UK are included in Part 4 (Specific Pollutants) of the above Directions. It should be noted that the UK Technical Advisory Group on the Water Framework Directive (UKTAG) is preparing a series of specific pollutant technical reports for the WDF, Annex VIII list of chemicals. These proposed EQS will only be used by Hydrock once they have been adopted formally by DEFRA and the Environment Agency.

The basis for the DWS and EQS used in this report is the Environment Agency Chemical Standards Database (http://evidence.environment-agency.gov.uk/ChemicalStandards/home.aspx).

The WFD imposes a duty on the Agency to classify surface water and groundwater bodies and to ensure long-term improvement (where necessary) to achieve acceptable standards. This includes the chemical status of the water. A new set of criteria for groundwater quality classification has been published, in addition to DWS and EQS. These are Threshold Values (TV) for individual groundwater bodies (GWB). Each GWB has been identified by the Agency and specific TVs calculated based on the perceived risks to that GWB. Failure of a TV is an indicator of potential adverse impact in specific circumstances. These TVs are not intended to be applied to meet the 'prevent or limit' objective of the Agency (UKTAG September 2008) and they are not used by Hydrock in this report.

Generic criteria for contaminated soils which might result in groundwater contamination can be derived from generic assumptions using the Environment Agency (2006) Remedial Targets Methodology. A tiered approach is detailed in this document. In accordance with CLR 11, EQS and DWS can be used as generic water quality targets with respect to contamination of controlled waters.

It is clearly not cost-effective to analyse every water sample for all determinands. Hydrock has produced a default *de minimus* suite which includes a number of common water quality indicators plus a selection of the more common chemicals of potential concern, drawn from

the lists of Specific Pollutants and Priority Substances / Priority Hazardous Substances in Parts 4 and 5, respectively, of the 2009 Directions, plus additional common contaminants listed in the EPA-H1 Part 2 document, as being indicators of Good water quality under the terms of the Directive.

In addition to this, Hydrock will add to this list any chemicals identified as potential risks by reference to the conceptual site model.

Using the WQTs discussed above, the risks to groundwater and surface water from contaminants on site have been assessed according to the remedial targets methodology (RTM) prescribed by the Environment Agency (2006).

The Level 1 soil zone assessment considers whether the contaminant concentrations in the soil moisture are sufficient to impact the water receptor(s). It is a conservative model and compares soil pore water concentrations with the above criteria, taking no account of dilution, dispersion or attenuation. Pore water concentrations can be estimated by analysis of perched water samples, analysis of eluates produced in the laboratory by standard leaching of soil samples, or by calculation from physico-chemical properties of the substances. Calculation may be more appropriate for poorly soluble substances where retention times may not be long enough during the standard leaching tests to reach equilibrium. However, the Environment Agency (2009d) cautions that the use of published k_d values to calculate pore water concentrations "can lead to a conservative estimate of risk" and suggest that leaching tests may be designed for non-volatile organics using BS18772:2008.

The Level 2 groundwater assessment is applicable where groundwater quality data are available and compares these with the above criteria, again taking no account of dilution, dispersion or attenuation.

The remedial targets methodology also allows for more detailed assessment (soil Level 2, 3 or 4, or groundwater Level 3 or 4) for substances which fail the above-mentioned assessments. These are progressively more complex assessments and do take into account attenuation and/or dilution, as applicable to the conceptual exposure model. Such assessment is beyond the scope of this report.

Where more than one water quality target is available it is important to apply the one relevant to the critical receptor. The DWS apply to groundwater or to surface water used for abstraction and the EQS apply to surface water where the aquatic ecosystem is the receptor. EQS are available for *inland* surface waters (freshwater) and *other* surface waters (transitional and marine). Where the most appropriate water quality target cannot be determined with certainty, the lowest one is adopted in line with the precautionary principle.

For the purposes of this report, the site data are compared with the various targets as set out in Table 3.9

Scenario	Water Body Receptors	Secondary Receptors	Example Contaminant linkages	RTM Level and Samples Used (if Available)	Water Quality Targets
A	Groundwater.	Human health (abstraction).	Contaminants from site leach or seep into groundwater body and this is a (potential/actual) source of human consumption or a strategic resource.		DWS
A	Groundwater. Surface water.	Human health (abstraction).	Contaminants from site leach or seep into groundwater body and this feeds surface water by base flow. The surface water may be used for human consumption.	RTM Level 2 -	DWS
В	Groundwater. Surface water.	Aquatic ecosystem.	Contaminants from site leach or seep into groundwater body and this feeds surface water by base flow. The surface water may be an aquatic ecosystem.	Groundwater. RTM Level 1 - Soil leachate (including any calculated pore	EQS (inland)
С	Groundwater. Surface water.	Aquatic ecosystem.	Contaminants from site leach or seep into groundwater body and this feeds surface water by base flow. The surface water may be an aquatic ecosystem.	water concentrations) or pore water.	EQS (other)
D	Groundwater. Surface water.	Human health (abstraction). Aquatic ecosystem.	Contaminants from site leach or seep into groundwater body and this feeds surface water by base flow. The surface water may be used for human consumption and is an aquatic ecosystem.		DWS EQS (inland)
E	Surface water.	Human health (abstraction).	Contaminants from site leach or seep into surface water which may be used for human consumption.	RTM Level 1 - Soil leachate (including any	DWS
F	Surface water.	Aquatic ecosystem.	Contaminants from site leach or seep into surface water which may be an aquatic ecosystem.	calculated pore water concentrations) or pore water.	EQS (inland)
G	Surface water.	Aquatic ecosystem.	Contaminants from site leach or seep into surface water which may be an aquatic ecosystem.	Although not part of the RTM,	EQS (other)
Η	Surface water.	Human health (abstraction). Aquatic ecosystem.	Contaminants from site leach or seep into surface water which may be used for human consumption and is an aquatic ecosystem.	these scenarios are used to compare surface water data to the water quality targets.	DWS EQS (inland)

Table 3.9: Summary of Water Quality Risk Assessment Protocol

Notes:

Some EQS are water hardness dependent. This is measured either in the receiving water or in groundwater (if it is part of the pathway), or is estimated from national maps.

Inland waters EQS applicable to freshwater, other waters EQS applicable to marine or transitional waters.

Where both DWS and EQS are applicable, it is assumed that the EQS is for inland waters.

This table and the results of the assessment are considered as a first screening for potential risks of pollution of Controlled Waters. More specific requirements may be stipulated by the Environment Agency.

Note that in some instances the reporting limit (or detection limit) quoted by the laboratory may be greater than the water quality target that it is being assessed against. Where this is the case it is noted in the table. The current exercise is an initial screening assessment.

There are three main possible reasons for this. Firstly, it may be that the 'standard' method gives a relatively higher reporting limit, but that a lower one could be obtained using a more specialised technique. However, it would be disproportionately expensive to adopt the more costly specialist technique for this initial screening exercise. Secondly, it may be that the sample in question was not 'clean' because the matrix was contaminated by other substances which interfere with the analysis and so a less sensitive method has been used to protect the laboratory equipment. Thirdly, it may be that no method exists that can reach the required limit. Hydrock has contacted the Environment Agency's own National Laboratory Service and even they cannot reach low enough limits for several of the substances in the Hydrock default suite (Cr(VI), total cyanide, phenols and certain PAHs). Consequently, and depending on the particular chemicals, it may be possible to say for certainty because suitable techniques are not available. Methods are being continually updated and new ones may become available.

In some cases all samples are below the detection limit but above the water quality target. It is not possible to make any judgement about these. However, in other cases, even though the detection limit is greater than the water quality target, some sample results do exceed the target.

3.12.1 Petroleum Hydrocarbons in Water

With respect to hydrocarbons in water, the Water Supply (Water Quality) Regulations 1989 (as amended 1999) contained a prescribed concentration of 10 μ g/l for "dissolved or emulsified hydrocarbons (after extraction with petroleum ether); mineral oils". This was removed from the 2000 (consolidated 2007) Regulations. It was confirmed by email from the Drinking Water Inspectorate to Hydrock (1 November 2005) that dissolved hydrocarbons are no longer a prescribed substance under the Regulations. However, the 10 μ g/l limit did remain in the Private Drinking Water Regulations 1991 until their revision at the end of 2009.

In the absence of a prescribed concentration for drinking water, many Environment Agency officers continue to use the superseded value. This is perhaps because petroleum hydrocarbons are a hazardous substance (former List 1) under the WFD. There is, however, no clear UK policy on hydrocarbon contamination of controlled waters. This is partly because analyses for 'petroleum hydrocarbons' are fraught with complications concerning false positives, the results being method dependent and not restricted to petroleum products.

Guidance written by the Environment Agency on risk assessment of hydrocarbons in groundwater is dated 2009 but has never been officially released through the Agency's website, although the dissemination status of the document is given as publicly available. This gives a table of water quality targets for hydrocarbons and lists "TPH (dissolved or emulsified hydrocarbons)". No minimum reporting value (MRV) is quoted, the value that would equate to a *de minimus* concentration under the prevention objective. The target of 10 μ g/l is given and this is described as coming from the "Private Water Supply Regulations S! 1991 No. 2790 (due to be updated in 2009)". As mentioned above, the 2009 Regulations no longer list dissolved hydrocarbons.

Furthermore, the guidance also states that in cases where petroleum hydrocarbons have already entered the water, the Agency will regulate under its limit objective, rather than the prevention objective. This means that EQS or DWS will be appropriate. However, none exist.

In the absence of definitive guidance on petroleum hydrocarbons in water Hydrock recognises that it is not possible to provide EQS and so regulation with respect to aquatic ecosystems is impossible. However, it is possible to extend the use of DWS by calculating screening criteria for the speciated TPH fractions. This provides a rational, transparent and risk-based approach using established scientific principles, rather than simply adopting a withdrawn standard.

Whilst not strictly applicable to aquatic ecosystems, at least this approach can help inform the judgement as to the degree of degradation of a water body.

Accordingly Hydrock has calculated guidelines for drinking water quality based on the methodology proposed by the World Health Organisation (WHO, 2005). This is based on an adult consuming 2 litres of water per day. Whereas the WHO document assumes a body weight of 60kg, Hydrock has assumed 70kg in keeping with the UK Contaminated Land CLEA methodology.

A conservative allocation of 10% of the oral Tolerable Daily Intake (TDI) has been attributed to intake from drinking water. It is noted by the WHO (2005) that exposure from other sources would be expected to be very small and that it would be possible to allocate a greater percentage to drinking water if required. In other words, this approach is very conservative and is appropriate as an initial screening value and allows for potential additive toxicity and simultaneous exposure from other sources.

The TDIs used are the same as those used in the derivation of soil GACs and are listed in Table 3.10 along with the calculated health-based water quality targets for drinking water. Note, however, that the Environment Agency (2009d) states that when considering carbon bands, one does not know the range of toxicities and health effects of the individual chemicals, and it is precautionary to assume that the toxicological effects are additive when setting water quality targets even though the toxic endpoints and modes of action might in reality be quite different. The recommendation is to adopt a precautionary approach whereby the water quality target for each band is divided by the number of bands with detected concentrations.

Determinand	TDI (μg/kg/day)	Solubility (µg/l)	Water Quality Target (see note 1) (µg/l)	Notes
Ali EC5-EC6	5000	35900	17500 ¹	
Ali >EC6-EC8	5000	5370	17500 ¹	This concentration would be significantly above the solubility in water.
Ali >EC8-EC10	100	427	350 ¹	

Table 3.10: Calculated Water	Quality Targets for Petrol	eum Hydrocarbons in Dri	nking Water
	adding rangeto for rouro	Sam nyai oou sono in Bri	mang mator

Determinand	TDI (μg/kg/day)	Solubility (µg/l)	Water Quality Target (see note 1) (µg/l)	Notes
Ali >EC10-EC12	100	33.9	350 ¹	This concentration would be significantly above the solubility in water.
Ali >EC12-EC16	100	0.759	350 ¹	This concentration would be significantly above the solubility in water.
Ali >EC16-EC44	2000	0.00254	7000 ¹	This concentration would be significantly above the solubility in water.
Aro EC5-EC7	223	1780000	1 ¹	Based on the TDI for toluene as recommended by Environment Agency (2005) P5-080/TR3 gives 780. In reality the UK DWS for benzene = 1 takes precedence.
Aro >EC7-EC8	223	590000	700 ¹	Calculated as 780, WHO DWS = 700 takes precedence.
Aro >EC8-EC10	40	64600	140 ¹	
Aro >EC10- EC12	40	24500	140 ¹	
Aro > EC12- EC16	40	5750	140 ¹	
Aro >EC16- EC21	30	653	105 ¹	
Aro >EC21- EC44	30	6.61	105 ¹	This concentration would be significantly above the solubility in water.
Benzene	n/a	1780000	1	Calculation not possible as non-threshold substance, UK DWS = 1 takes precedence.
Toluene	223	590000	700	Calculated as 780, WHO DWS = 700 takes precedence.
Ethylbenzene	100	180000	300	Calculated as 350, WHO DWS = 300 takes precedence.
Xylene	180	200000	500	Calculated as 630, WHO DWS = 500 takes precedence.
MTBE	300	48000000	15	Calculated as 1050 so the odour threshold = 15 is adopted.
Note 1: The value to	be used in a ris	k assessment (for carbon bands) is the value in the table divided by the number of

Note 1: The value to be used in a risk assessment (for carbon bands) is the value in the table divided by the number of bands with detected concentrations.

Last updated 29/06/10

In instances where a simple 'total' TPH is reported for water samples this should be considered indicative only. This is particularly the case if groundwater or surface water samples were not available and an indication of pore water quality has been derived by subjecting soil samples to a standard leaching procedure or calculation.

Where petroleum hydrocarbon contamination of Controlled Waters is suspected, Hydrock recommends that discussion with the Environment Agency is entered into at the earliest opportunity.

3.13 Statistical Tests of Soil Contamination Results

As discussed above, the sample analyses are divided into representative data sets for the assessment, based on the conceptual site model, and are referred to as 'averaging areas'. In this case it has been chosen to characterize materials that are likely to form the ground cover in critical receptor areas (e.g. gardens), on a material by material basis. The critical

part of the soil column is the upper metre in terms of contact with end users of a development site.

Under the **land use planning system** where the aim is to demonstrate 'suitability for use' the key question will usually be "can we say confidently that the level of contamination of this land is low relative to some appropriate measure of risk, sometimes referred to as the critical concentration?" The critical concentration can be, for example, the relevant GAC.

It is necessary to demonstrate that (for each contaminant) the mean concentration on the site is **below** the critical concentration. The true mean concentration of a contaminant is not known because all the site soil has not been tested. An estimation of the true mean can be obtained from the samples tested during the investigation. The greater the number of samples tested, the closer the mean of these values is to the true mean.

In practice, this involves calculation of a quantity known as the 95th Upper Confidence Limit (UCL) of the true population mean, also known as the US_{95} . This is the estimate of the true mean at a 95% level of confidence (i.e. there is a 95% probability that the true mean will not be greater than this, given the values obtained from the investigation sample testing).

The statistical test that is carried out, therefore, is used to demonstrate that there is a 95% probability that the true mean falls below critical concentration (typically the GAC in a screening exercise).

In statistical language, a **null hypothesis** is stated; that the level of contamination is the same as, or higher than, the critical concentration. The **alternative hypothesis** is that the level of contamination is lower than the critical concentration. The statistical test is used to decide whether or not the null hypothesis is rejected.

If it is rejected, the assessor can conclude that the alternative hypothesis is more likely to be true, i.e. that contaminant concentrations are low relative to the critical concentration and that, potentially, the land is suitable for use. Conversely, if the null hypothesis is not rejected, the assessor should conclude that contaminant concentrations may be the same as, or higher than, the critical concentration and further measures may be needed.

It should be noted that a similar, but opposite, set of propositions applies in the case of a potential Part 2A determination where the level of contamination must be higher than some appropriate level of risk (critical concentration) (e.g. that indicative of SPOSH). In this case, however, a lower standard of proof may be accepted and the guidance suggests that if the statistical test of significance at the 95% confidence level does not indicate rejection of the null hypothesis, then the test should be repeated at the 51% level to see if there is evidence to suggest the null hypothesis be rejected on the balance of probabilities.

A useful summary of the methodology is provided by CIEH & CL:AIRE (May 2008), which forms the basis for the approach adopted by Hydrock, and is described below.

Firstly, the data set is assessed for outliers and normality. This is mainly a visual exercise rather than following a particular statistical method. The reason for this approach was that the former guidance provided in CLR7 had the potential for misuse and that is why it was withdrawn. It is considered that a reasoned option is preferable to blindly following a particular test when it comes to outliers and normality.

Two graphs are considered, the data frequency histogram with a normal 'bell curve' for comparison and a quantlie-quantile (q-q) plot. The closer the data points lie to the 45° line, the closer they are to a normal distribution. Kinks in the q-q plot are indicative of more than one data set. Individual points away from the 45° line are indicative of outliers.

Additional evidence of outliers is obtained through a simple method of robust statistics advocated by the Royal Society of Chemistry (2001) and others. The measure of the mean is taken to be the median value because this is less susceptible to outliers and non-normal data sets. A value known as the mean absolute deviation (MAD) is calculated and from this can be calculated a robust standard deviation estimate by multiplying by 1.483.

A z-score can then be calculated, which is the absolute value of the data value minus the median, divided by the robust standard deviation. This is then compared with a critical value which, if exceeded, suggests a possible outlier. The critical value represents the number of standard deviations from the mean (or in this case the median). A critical value of 3 to 3.5 is generally considered appropriate. The attraction of this approach is that it is a robust, non-parametric method suitable for all data sets. It is not considered as definitive, but merely a tool to aid decision making.

If a potential outlier is identified it could be a laboratory or typographic error. If this is not the case it could be representative of a different contaminative incident and, therefore, be a hot-spot. However, it could also be simply the result of heterogeneous ground conditions and a relatively low number of sampling points. The initial review of the data is then coupled to a knowledge of the conceptual site model before an outlier is removed from the data set. A good reason is required to justify the removal of outliers and this will be reported in the text.

The second stage of the assessment is to carry out the statistical test as described previously. Two alternative methods are highlighted in the CIEH/CL:AIRE document. The one-sample t-test is appropriate for normally distributed data (it is a parametric test) but is not sensitive to moderate departures from normality. The Chebychev Theorem is a non-parametric test which is suitable for all data distributions. It is a less powerful test (statistically) and gives a more cautious result than the t-test because there is less certainty about the shape of the distribution.

If there is significant departure from normality, perhaps because outliers are still included, the t-test will be chosen. Otherwise, the Chebychev Theorem will be used. The chosen method is applied and the outcome recorded with respect to whether or not the null hypothesis is rejected and the site is potentially suitable for use.

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When considering potential Part 2A sites, updated guidance published by Barnes *et al* (2010) recommends the t-test for all data sets (unless the data are negatively skewed, something these authors have never seen in contaminated land data sets).

Please note that under certain circumstances a 'divided by 0' error can occur in the spreadsheets used in the statistical analyses. This happens when all the data points are the same integer value, for example where all results are <3 mg/kg and they have been assumed to be 3 mg/kg. To prevent this error, one of the results can be altered by a small amount (e.g. 3 becomes 2.99999). This allows the statistical tests to be carried out but makes no difference to the outcome. However, it does mean that the q-q and histogram plots show a spurious point, which should be ignored.

3.13.1 Note on Clustered Data Sets

The assumption behind the statistical tests is that each sample represents an equal fraction of the averaging area (Nathanail, 2004). If the data are clustered, i.e. the sampling points are not equally spaced, the calculated US_{95} would be too high if targeted sampling has taken place around suspected high concentration areas to determine the extent of the high contamination. Conversely, the calculated US_{95} would be too low if there is a high density of sampling in an area of low contaminant concentration.

The sampling pattern used in this report has been reviewed to determine if clustering of data points is likely to affect the statistical tests significantly. In cases where the area represented by each sample is judged to be similar, the tests have been carried out without modification. The error in this approach is likely to be conservative to human health because the Hydrock approach to targeted sampling is more likely to produce more closely spaced higher concentrations than more closely spaced lower concentrations.

Erring on the conservative side is, however, counter productive when it would indicate unnecessary remediation, i.e. remediation triggered by a US₉₅ which is skewed by clustered data. This is taken into consideration in the risk evaluation part of the risk assessment exercise and can take the form of professional judgement, the modification of the averaging area datasets to decluster them, or the weighting of sample results to decluster the data set. The latter method involves weighting the measured concentrations according the proportion of the area they represent, giving greater weight to samples representative of a larger area.

3.13.2 Statistical Tests and Risk to Controlled Waters

Where only a few water quality tests are available, the maximum concentrations are compared with the standards because the 95 percentile will be close to the maximum value. However, where a larger population is available, the 95 percentile is compared with the standards, as recommended by the Environment Agency.

3.14 Ground Gas Risk Assessment

The risks associated with the ground gases methane (CH_4) and carbon dioxide (CO_2) are assessed using BS 8485:2007 and guidelines from CIRIA (Wilson *et al* 2007) and the NHBC (Boyle and Witherington 2007).

In the above guidance, 'Situation B' is defined as the specific development of low-rise (1 or 2 storey) housing with beam and block floors, vented sub-floor void and gardens. Initial risk classification can be made according to NHBC Table 8.1. This determines the appropriate risk strategy for protection, including the need to progress to generic quantitative risk assessment (GQRA). Even where no risk assessment is recommended by this table, one may be carried out if so desired. The GQRA is known as the 'NHBC traffic light classification' as it uses red, amber and green designations to portray levels of risk.

'Situation A' covers all other forms of development. This uses a modified version of the Wilson and Card (1999) methodology.

The idealised frequency of monitoring is suggested in CIRIA Tables 5.5a and 5.5b. These tables are adapted from Wilson and Haines (2005) Table 3 which gives examples of ground conditions with the various gas generation potentials, ranging from inert Made Ground (very low potential) to post 1960s domestic landfill (very high potential).

The report does not constitute a design for gas protection measures, but lists the recommendations given by the above-mentioned guidance for the particular "Situation" considered relevant. For information, BS 8485:2007, Table 3 is reproduced herein as Table 3.11.

Protection Element/System	m	Score	Comments
a) Venting/dilution (see B	S 8485, Annex A)		
Passive sub floor ventilation (venting layer	Very good performance.	2.5	Ventilation performance in accordance with Annex A.
can be a clear void or formed using gravel, geocomposites, polystyrene void formers, etc.). ^{A)}	Good performance.	1	If passive ventilation is poor this is generally unacceptable and some form of active system will be required.
Subfloor ventilation with act abstraction/pressurization (void or formed using gravel polystyrene void formers, et	venting layer can be a clear , geocomposites,	2.5	There have to be robust management systems in place to ensure the continued maintenance of any ventilation system. Active ventilation can always be designed to meet good performance. Mechanically assisted systems come in two main forms: extraction and positive pressurization.

Protection Element/Systemeter	em	Score	Comments	
Ventilated car park (basement or undercroft).		4	Assumes car park is vented to deal with car exhaust fumes, designed to Building Regulations Document F and IStructE guidance (design recommendations for multistorey and underground car parks, 3rd ed, 2002).	
b) Barriers				
Block and beam floor slab.		0	It is good practice to install ventilation in all foundation systems to effect pressure relief as a minimum. Breaches in floor slabs such as joints have to be effectively sealed against gas ingress in	
Reinforced concrete ground bearing floor slab.		0.5		
Reinforced concrete ground bearing foundation raft with limited service penetrations that are cast into slab.		1.5		
Reinforced concrete cast in situ suspended slab with minimal service penetrations and water bars around all slab penetrations and at joints.		1.5	order to maintain these performances.	
Fully tanked basement.		2]	
c) Membranes				
Taped and sealed membrane to reasonable levels of workmanship/in line with current good practice with verification. ^{B), C)}		0.5	The performance of membranes is heavily dependent on the quality and design of the installation, resistance to damage after	
Proprietary gas resistant membrane to reasonable levels of workmanship/in line with current good practice under independent inspection (CQA). ^{B), C)}		1	installation and the integrity of joints.	
Proprietary gas resistant membrane installed to reasonable levels of workmanship/in line with current good practice under CQA with integrity testing and independent verification.		2		
d) Monitoring and detect	ion (not applicable to non-m	nanaged p	roperty, or in isolation)	
Intermittent monitoring usin	ermittent monitoring using hand held equipment 0.5 Where fitted, permanent mon		Where fitted, permanent monitoring systems	
Permanent monitoring and alarm system. A)	Installed in the underfloor venting/dilution system.	2	ought to be installed in the underfloor venting/dilution system in the first instance but can also be provided within the occupied space as a fail safe.	
	Installed in the building.	1		
e) Pathway intervention				
Pathway intervention.		-	This can consist of site protection measures for off-site or on-site sources (see Annex A).	
	of materials might well rely on fac nsure that the chosen combinatio		s construction method and the risk of damage after propriate level of protection.	
 B) If a 1200 g DPM material (Johnson 2001), being tap C) Polymeric Materials >120 	bed and sealed to all penetrations 0 g can be used to improve confid	nould be inst lence in the	for post installation verification. alled according to BRE 212 (Hartless 1991)]/BRE 414 barrier. Remember that their gas resistance is little sical properties mean that they are more robust and	

4.0 WATER SUPPLY PIPES

The current guidance on selection of materials for water supply pipes to be laid in contaminated land is contained in UKWIR Report 10/WM/03/02 (re-issued 2010) which sets out in Table 3.1 of that document threshold values for a selection of organic contaminants that may have a detrimental effect on pipes and fittings. The contaminants are divided into a number of 'parameter groups'. Also included are threshold values for certain parameters that could cause corrosion of metal pipes. This guidance supersedes the former WRAS guidance (October 2002).

Some water suppliers (e.g. Anglian, United Utilities, Wessex and Yorkshire) specify their own soil threshold values and material design requirements, however these are currently not nationally recognised.

The UKWIR guidance recommends a *mandatory* analytical suite in its Table G1 comprising five groups of substances as follows: extended VOC suite, BTEX & MTBE, extended SVOC suite, phenols, cresols and chlorinated phenols, mineral oil C11-C20, mineral oil C21-C40 and corrosive suite. Other groups and sub-groups are recommended on a site-specific basis: ethers, nitrobenzene, ketones, aldehydes and amines.

The Association of Geotechnical and Geoenvironmental Specialists (AGS, 9 July 2009) stated that the former WRAS guidance was out of date and should be withdrawn and that the internal guidance adopted by water companies is inconsistent in some respects. Unfortunately, the replacement UKWIR document also contains a number of technical errors and inconsistencies and to date has not been universally accepted. For example, it requires a desk study, but then gives a mandatory list of determinands that would negate the need for a desk study. Also, a number of substances are listed under the wrong groups in Table G1. The AGS continues to have concern.

The UKWIR suggested methodology includes a soil vapour survey in cored boreholes along the pipeline routes, the construction of sampling boreholes where water is detected within 1 m of the base of a proposed pipe trench and analysis of soil samples over a 30 m wide corridor along each pipe run. The sampling plan is to be agreed with the water company beforehand. There are significant cost and time implications associated with this approach and so it has **not** been followed as part of this report.

Note that the use of barrier pipe (PE-AI-PE) is applicable for all brownfield sites according to the UKWIR guidelines. Further work may be required if other types are contemplated.

As a minimum, the findings of this investigation can been compared to the threshold values in UKWIR Table 3.1 as far as is practicable and with the modifications shown in Table 4.1 (and ignoring obvious errors in Table G1) to give an indication of the possible restrictions to the use of plastic water pipes.

Parameter Group from UKWIR Table 3.1	Hydrock Comments	Trigger Value (mg/kg)	
		PE	PVC
1 – Extended VOC suite by purge & trap or head-space and GC-MS with TIC (total VOC concentration excluding BTEX and MTBE).	Sum of VOC excluding BTEX & MTBE from this report (if available).	0.5	0.125
1a – BTEX & MTBE (total concentration).	Sum of BTEX and MTBE from this report (if available).	0.1	0.03
	Sum of SVOC from this report excluding substances in 2a to 2f.		
2 – Extended SVOC suite by purge & trap or head-space and GS-MS with TIC (aliphatic and aromatic C5-C10) (total SVOC concentration excluding 2a to 2f listed below).	Also excluding petroleum hydrocarbons aliphatic and aromatic C5-C10 listed by UKWIR because hese are VOCs not SVOCs and it is presumed to be an error. These will be included in the VOC total automatically.		1.4
2e – Phenols (total concentration).	Sum of phenols from this report (if available).	2	0.4
2f – Cresols and chlorinated phenols (total concentration).	Sum of cresols and chlorinated phenols from this report (if available).	2	0.04
3 – Mineral oil C11-C20 (total concentration).	Sum of TPH fractions >EC10-EC21 from this report (if available).	10	No limit
4 – Mineral oil C21-C40 (total concentration).	Sum of TPH fractions >EC21-EC35 from this report (if available).	500	No limit
2a – Ethers (if site history suggests as a potential hazard) (total concentration).	Sum of ethers from this report (if available).	0.5	1
2b – Nitrobenzene.	Nitrobenzene (if available).	0.5	0.4
2c – Ketones (total concentration).	Sum of ketones from this report (if available).	0.5	0.02
2d – Aldehydes (total concentration).	Sum of aldehydes from this report (if available).	0.5	0.02
6 – Amines (total concentration).	The UKWIR report says the presence of amines precludes the use of PE. Since a less than detection limit value is not necessarily zero, logically PE cannot be used on site where the desk study leads to a suspicion of there being amines in the ground.	Fail	No limit

In view of the lack of clear and unambiguous guidance it is **strongly recommended** that site-specific approval of the materials for underground pipes to be used for water supply be obtained from the water company that will be supplying this site and/or adopting the pipework.

5.0 FLOOD RISK

The following additional information concerns the background to flood risk mentioned in the report. Guidance is given in the document *Technical Guidance to the National Planning Policy Framework* (DCLG March 2012) which retains key elements from the withdrawn Planning Policy Statement 25.

The Environment Agency flood maps are divided into Flood Zones, as follows.

- Flood Zone 1 is land outside the extent of extreme flooding and the annual risk is less than 1:1000, low probability (depicted as white on the web-based map).
- Flood Zone 2 is land unlikely to flood except in extreme conditions if no defences are present and the annual risk is between 1:100 and 1:1000 (for rivers) or 1:200 and 1:1000 (for the sea), medium probability (depicted as light blue on the web-based map).
- Flood Zone 3 is land within the floodplain at risk of flooding if no defences are present and the annual risk is greater than or equal to 1:100 (for rivers) or 1:200 (for the sea), high probability (depicted as dark blue on the web-based map).

The Agency flood maps also define the risk of flooding: as 'low' (\leq 1:200), 'moderate' (>1:200 to \leq 1:75) or 'significant' (>1:75), which are not the same divisions as those in the guidance mentioned above. Note that the published flood map only relates to flooding from rivers, estuaries and the sea and does not include other potential sources such as surface water, groundwater, sewers, canals and reservoirs. Note also that the presence on the map of flood defences, or areas benefiting from flood defences, should not be taken to imply that a proposed development in these areas is acceptable.

The **Environment Agency in England** has issued Flood Risk Standing Advice. However, this is to be reviewed following the publication of the NPPF (see http://www.environment-agency.gov.uk/research/planning/33098.aspx for updates and details).

The flood map mentioned above can be accessed at the Agency's website.

The Technical Guidance states:

 Within Flood Zone 1 all uses of land are appropriate. For development proposals on sites comprising one hectare or above, the vulnerability to flooding from other sources as well as from river and sea flooding; and the potential to increase flood risk elsewhere through the addition of hard surfaces and the effect of the new development on surface water run-off, should be incorporated in a flood risk assessment (FRA) to accompany the planning application. This need only be brief unless the factors above or other local considerations require particular attention. For development proposals less than one hectare no flood risk assessment (FRA) is required.

- Within Flood Zone 2, water-compatible, less vulnerable and more vulnerable uses of land and essential infrastructure (as defined in Technical Guidance, Table 2) are appropriate in this zone. The Sequential Test of PPS25 is required and must be passed and for highly vulnerable uses in Table 2 the Exception Test must be applied and passed also. All development proposals in this zone should be accompanied by a flood risk assessment (FRA).
- Flood Zone 3 is sub-divided into 3a and 3b, but these are not distinguished on the published maps. Flood Zone 3a is land having an annual probability of flooding of >1:100 (from rivers) or >1:200 (from the sea). The water-compatible and less vulnerable uses of land (as defined in Technical Guidance, Table 2) are appropriate in this zone. The highly vulnerable uses in Table 2 should not be permitted in this zone. The Sequential Test is required and must be passed and for the more vulnerable and essential infrastructure uses in Table 2 the Exception Test must be applied and passed also. Essential infrastructure permitted in this zone should be designed and constructed to remain operational and safe for users in times of flood. All development proposals in this zone should be accompanied by a flood risk assessment (FRA).

Flood Zone 3b is known as the 'functional floodplain' and comprises land where water has to flow or be stored in times of flood and should be identified on Strategic Flood Risk Assessments (SFRA) undertaken by the Local Planning Authority. Such land is defined as land which would flood with an annual probability of 1:20 or greater, or is *designed* to flood in an extreme (1:1000) flood, or at another probability to be agreed between the Local Planning Authority and the Environment Agency, including water conveyance routes). Only the water-compatible uses and the essential infrastructure (as defined in Technical Guidance, Table 2) that has to be there should be permitted in this zone. It should be designed and constructed to: remain operational and safe for users in times of flood; result in no net loss of floodplain storage; not impede water flows; and not increase flood risk elsewhere. The Sequential Test is required and must be passed and for essential infrastructure the Exception Test must be applied and passed also. All development proposals in this zone should be accompanied by a FRA.

The **Environment Agency Wales** flood map is not used for planning purposes (only to provide information on flood risk and to raise awareness). Development advice with respect to flooding is provided by the Welsh Assembly Government (July 2004) Technical Advice Note 15 (TAN15) and the accompanying development advice maps. An interactive map is available from the WAG web site.

The development advice map containing three zones (A, B and C with subdivision into C1 and C2) should be used to trigger the appropriate planning tests.

- Zone A is considered to be at little or no risk of fluvial or tidal/coastal flooding. The justification test (TAN15, Section 6) is not applicable and there is no need to consider flood risk further. This equates to Flood Zone 1 on the Agency maps.
- Zone B is land known to have been flooded in the past evidenced by sedimentary deposits. As part of a precautionary approach site levels should be checked against the extreme (1:1000) flood level. If site levels are greater than the flood levels used to define adjacent extreme flood outline there is no need to consider flood risk further. This land within Flood Zone 1 of the Agency maps but close to Flood Zone 2 or 3.

Zone C is based on the Environment Agency Wales extreme flood outline, equal to or greater than 1:1000 (river, tidal or coastal) and equates to Flood Zones 2 and 3 on the Agency map. Flooding issues should be considered as an integral part of decision making by the application of the justification test (TAN15, Section 6) including assessment of consequences (TAN15, Section 7) is required. Sub-division C1 is land in the floodplain which are developed and served by significant infrastructure, including flood defences. Development can take place subject to application of the justification test, including acceptability of consequences. Sub-division C2 is land in the floodplain without significant flood defence infrastructure. Only less vulnerable development should be considered subject to application of the justification test, including acceptability of consequences. Emergency services and highly vulnerable development should not be considered. The categories of land use are defined in TAN15, Figure 2.

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Appendix E

CONTAMINATION TEST RESULTS & STATISTICAL ANALYSIS

This appendix may not be included in the printed report to reduce the document size. It is presented in the PDF version of the report on the CD enclosed with the printed report.



Depot Road Newmarket CB8 0AL Tel: 01638 606070

Hydrock Consultants Over Court Barns Over Lane Almondsbury, Bristol BS32 4DF

FAO Rob Hooker 14 December 2012

Dear Rob Hooker

Test Report Number217776Your Project ReferenceBanbury

Please find enclosed the results of analysis for the samples received 10 December 2012.

All soil samples will be retained for a period of one month and all water samples will be retained for 7 days following the date of the test report. Should you require an extended retention period then please detail your requirements in an email to customerservices@chemtest.co.uk. Please be aware that charges may be applicable for extended sample storage.

If you require any further assistance, please do not hesitate to contact the Customer Services team.

Yours sincerely



Darrell Hall, Director





Notes to accompany report:

- The sign < means 'less than'
- Tests marked 'U' hold UKAS accreditation
- Tests marked 'M' hold MCertS (and UKAS) accreditation
- Tests marked 'N' do not currently hold UKAS accreditation
- Tests marked 'S' were subcontracted to an approved laboratory
- n/e means 'not evaluated'
- i/s means 'insufficient sample'
- u/s means 'unsuitable sample
- Comments or interpretations are beyond the scope of UKAS accreditation
- The results relate only to the items tested
- All results are expressed on a dry weight basis
- The following tests were analysed on samples as received and the results subsequently corrected to a dry weight basis TPH, BTEX, VOCs, SVOCs, PCBs, phenols
 - For all other tests the samples were dried at < 37°C prior to analysis
- Uncertainties of measurement for the determinands tested are available upon request
- None of the test results included in this report have been recovery corrected

Test Report 217776 Cover Sheet



Depot Road Newmarket CB8 0AL Tel: 01638 606070

Hydrock Consultants Over Court Barns Over Lane Almondsbury, Bristol BS32 4DF

FAO Rob Hooker 14 December 2012

Dear Rob Hooker

Test Report Number217776Your Project ReferenceBanbury

Please find enclosed the results of analysis for the samples received 10 December 2012.

If you require any further assistance, please do not hesitate to contact the Customer Services team.

Yours sincerely

Darrell Hall, Director



Notes to accompany report:

- The in-house procedure is employed to identify materials and fibres in soils
- The sample is examined by stereo-binocular and polarised light microscopy
- Sample size is reduced by coning and quartering to obtain a representative sub-sample if necessary
- The bulk identification is in accordance with the requirements of the analyst guide (HSG 248)
- Samples associated with asbestos are retained for six months
- The results relate only to the items tested as supplied by the client
 - Comments or interpretations are beyond the scope of UKAS accreditation



Test Report 217776 Cover Sheet

FAO Rob Hooker

Login Batch No: 217776

Qualitative Results

Quantative N	counto				SOP 2190
				АСМ Туре	Asbestos Identification
Chemtest ID	Sample ID	Sample Desc	Depth (m)		
AI05105	7	HTP02	0.20	-	No Asbestos Detected
AI05108	12	HTP03	0.50	-	No Asbestos Detected
AI05109	16	HTP04	0.10	-	No Asbestos Detected
AI05112	23	HTP05	0.50	-	No Asbestos Detected
AI05113	28	HTP06	0.10	-	No Asbestos Detected
AI05115	33	HTP07	0.50	-	No Asbestos Detected
AI05117	41	HTP09	0.20	-	No Asbestos Detected
AI05118	47	HTP10	0.50	-	No Asbestos Detected
AI05121	116	HTP13	0.20	-	No Asbestos Detected
AI05123	101	HTP15	0.25	-	No Asbestos Detected
AI05125	109	HTP17	0.15	-	No Asbestos Detected
AI05127	129	HTP20	0.15	-	No Asbestos Detected
AI05129	137	HTP22	0.20	-	No Asbestos Detected
AI05131	76	HTP24	0.20	-	No Asbestos Detected
AI05132	72	HTP25	0.15	-	No Asbestos Detected
AI05133	68	HTP26	0.10	-	No Asbestos Detected
AI05135	87	HTP29	0.15	-	No Asbestos Detected
AI05137	64	HTP33	0.25	-	No Asbestos Detected
AI05140	90	HTP39	0.20	-	No Asbestos Detected
AI05141	97	HTP40	0.20	-	No Asbestos Detected

The detection limit for this method is 0.001%

Signed

Signed

Pauline Hellier Asbestos Analyst



Results of analysis of 20 samples

received 10 December 2012

Banbury



Report Date 14 December 2012

LABORATORY TEST REPORT



Results of analysis of 4 samples received 10 December 2012

FAO Rob Hooker

Report Date 14 December 2012

Banbury

Login E	Batch No					217	776	
Chemte	est LIMS ID				AI05256	AI05257	AI05258	AI05259
Sample	ID				HTP02	HTP05	HTP11	HTP35
Sample	No				8	22	51	57
Samplin	ng Date				Not Provided	Not Provided	Not Provided	Not Provided
Depth					0.50m	0.20m	0.50m	0.60m
Matrix					LEACHATE	LEACHATE	LEACHATE	LEACHATE
SOP↓	Determinand↓	CAS No↓	Units↓	*				
1010	рН	PH		U	8.2	8.0	8.0	7.9
1020	Electrical Conductivity	EC	µS cm-¹	U	220	240	36	80
1215	Bromate	15541454	mg l-1	Ν	<0.01	<0.01	<0.01	<0.01
1220	Chloride	16887006	mg l-1	U	2.5	2.7	2.9	1.4
	Fluoride	16984488	mg l-1	U	0.16	0.25	0.34	0.62
	Ammonium	14798039	mg l-1	U	0.20	0.20	0.37	0.27
	Nitrite	14797650	mg l-1	U	0.12	0.44	0.16	0.19
	Nitrate	14797558	mg l-1	U	0.64	5.7	1.4	2.3
1300	Cyanide (free) Low-Level	57125	mg l-1	Ν	< 0.005	<0.005	<0.005	< 0.005
	Cyanide (total) Low-Level	57125	mg l-1	Ν	<0.005	<0.005	<0.005	<0.005
1470	Iron (dissolved)	7439896	µg l-¹	Ν	590	100	1200	220
1270	Hardness	HARD_TOT	mg CaCO3 I-1	U	61	73	28	280
1415	Sodium	7440235	mg l-1	U	2.3	1.6	2.3	5.3
1220	Sulfate	14808798	mg l-1	U	6.8	3.6	6.7	1.6
1430	Aluminium	7429905	µg l-¹	Ν	1200	30	770	170
1450	Silver	7440224	µg l-1	Ν	<0.5	<0.5	<0.5	<0.5
	Arsenic	7440382	µg l-¹	U	1.6	<1.0	<1.0	<1.0
	Boron	7440428	µg l−¹	U	<20	<20	<20	<20
	Barium	7440393	µg l-¹	U	6.2	33	<5.0	<5.0
	Cadmium	7440439	µg l-¹	U	<0.080	<0.080	<0.080	<0.080
	Cobalt	7440484	µg l-¹	U	<1.0	<1.0	<1.0	<1.0
	Chromium	7440473	µg l-¹	U	<1.0	<1.0	<1.0	<1.0
	Copper	7440508	µg l-¹	U	<1.0	6.3	1.7	<1.0

¹No sampling date was specified, stability times for this analyte may have been exceeded and these results may be compromised. The accreditation for these results remains unaffected.

All tests undertaken between 10/12/2012 and 14/12/2012

* Accreditation status

Column page 1 Report page 1 of 6 LIMS sample ID range Al05103 to Al05259

LABORATORY TEST REPORT



Results of analysis of 4 samples received 10 December 2012

Banbury

Report Date 14 December 2012

FAO Rob Hooker

					217776						
					AI05256	AI05257	AI05258	AI05259			
					HTP02	HTP05	HTP11	HTP35			
					8	22	51	57			
					Not Provided	Not Provided	Not Provided	Not Provided			
					0.50m	0.20m	0.50m	0.60m			
					LEACHATE	LEACHATE	LEACHATE	LEACHATE			
450	Managana	7439965			00	-11.0	F 0	1.0			
450	Manganese		µg l-1	U	29	<1.0	5.8	1.2			
	Molybdenum	7439987	µg l-1	U	<1.0	19	<1.0	<1.0			
	Nickel	7440020	µg l-1	U	<1.0	<1.0	<1.0	<1.0			
	Lead	7439921	µg l-¹	U	<1.0	2.0	<1.0	<1.0			
	Antimony	7440364	µg l-¹	U	<1.0	<1.0	<1.0	<1.0			
	Selenium	7782492	µg l-¹	U	1.8	<1.0	<1.0	<1.0			
	Tin	7440315	µg l₋¹	U	2.2	<1.0	<1.0	<1.0			
	Vanadium	7440622	µg l-¹	U	2.6	<1.0	1.8	<1.0			
	Zinc (dissolved)	7440666	µg l-¹	U	3.4	3.8	4.2	<1.0			
	Zinc	7440666	µg l-¹	U	3.5	3.8	4.2	<1.0			
460	Mercury Low Level	7439976	µg l₋¹	U	0.022	<0.01	<0.01	< 0.01			
490	Chromium (trivalent)	16065831	µg l₋¹	Ν	<20	<20	<20	<20			
	Chromium (hexavalent)	18540299	µg l-¹	U	<20 ¹	<20 ¹	<20 ¹	<20 ¹			
700	Naphthalene	91203	µg l-1	Ν	2.6	3.3	2.2	< 0.01			
	Acenaphthylene	208968	µg l-1	N	0.5	0.6	0.6	0.8			
	Acenaphthene	83329	µg l-1	Ν	1.8	1.9	0.7	1			
	Fluorene	86737	µg l-1	N	94	69	<0.01	< 0.01			
	Phenanthrene	85018	µg l-1	N	2	1.6	<0.01	<0.01			
	Anthracene	120127	µg l-1	N	<0.01	<0.01	<0.01	<0.01			
	Fluoranthene	206440	µg l-1	Ν	< 0.01	0.6	<0.01	1			
	Pyrene	129000	µg l-1	Ν	<0.01	1.3	<0.01	1.6			
	Benzo[a]anthracene	56553	µg l-1	Ν	<0.01	<0.01	<0.01	< 0.01			
	Chrysene	218019	μg l-1	N	<0.01	< 0.01	< 0.01	< 0.01			
	Benzo[b]fluoranthene	205992	μg l-1	N	<0.01	1.2	<0.01	< 0.01			

¹No sampling date was specified, stability times for this analyte may have been exceeded and these results may be compromised. The accreditation for these results remains unaffected.

All tests undertaken between 10/12/2012 and 14/12/2012

* Accreditation status

Column page 1 Report page 2 of 6 LIMS sample ID range AI05103 to AI05259

LABORATORY TEST REPORT



Results of analysis of 4 samples received 10 December 2012

Banbury

Report Date 14 December 2012

FAO Rob Hooker

					217776					
					AI05256	AI05257	AI05258	AI05259		
					HTP02	HTP05	HTP11	HTP35		
					8	22	51	57		
					Not Provided	Not Provided	Not Provided	Not Provided		
					0.50m	0.20m	0.50m	0.60m		
					LEACHATE	LEACHATE	LEACHATE	LEACHATE		
1700	Benzo[k]fluoranthene	207089	µg l-¹	N	<0.01	<0.01	<0.01	<0.01		
1700	Benzo[a]pyrene	50328	μg I-1	N	8.9	6.7	5.2	7.1		
	Dibenzo[a,h]anthracene	53703	μg I-1	N	2.6	2.6	1.6	<0.01		
	Indeno[1,2,3-cd]pyrene	193395	µg I-1	N	<0.01	<0.01	<0.01	<0.01		
	Benzo[g,h,i]perylene	193395	µg I-1	N	<0.01	<0.01	<0.01	<0.01		
	Total (of 16) PAHs	191242	μg I-1	N				11		
1000	· · ·	88857		N	110 <0.0002	89 <0.0002	10	<0.0002		
1900	2-sec-Butyl-4,6-dinitrophenol 4-Chloro-3-methylphenol	59507	mg l-1	N						
	71		mg l-1		< 0.0002	< 0.0002	< 0.0002	< 0.0002		
	2-Chlorophenol	95578 120832	mg l-1	N	< 0.0002	< 0.0002	< 0.0002	< 0.0002		
	2,4-Dichlorophenol		mg l-1	N	< 0.0002	< 0.0002	< 0.0002	< 0.0002		
	2,6-Dichlorophenol	87650	mg l-1	N	<0.0002	< 0.0002	<0.0002	< 0.0002		
	2,4-Dimethylphenol	105679	mg l-1	N	<0.0002	<0.0002	<0.0002	<0.0002		
	2,4-Dinitrophenol	51285	mg l-1	Ν	<0.0002	<0.0002	<0.0002	<0.0002		
	2-Methyl-4,6-dinitrophenol	534521	mg l-1	N	<0.0002	<0.0002	<0.0002	<0.0002		
	2-Methylphenol	95487	mg l-1	N	< 0.0002	< 0.0002	< 0.0002	< 0.0002		
	3-Methylphenol	108394	mg l-1	N	<0.0002	< 0.0002	<0.0002	< 0.0002		
	4-Methylphenol	106445	mg l-1	Ν	< 0.0002	< 0.0002	< 0.0002	< 0.0002		
	2-Nitrophenol	88755	mg l-1	N	< 0.0002	<0.0002	<0.0002	< 0.0002		
	4-Nitrophenol	100027	mg l-1	Ν	< 0.0002	<0.0002	< 0.0002	< 0.0002		
	Pentachlorophenol	87865	mg l-1	Ν	< 0.0002	< 0.0002	< 0.0002	< 0.0002		
	Phenol	108952	mg l-1	N	< 0.0002	< 0.0002	< 0.0002	< 0.0002		
	2,3,4,5-Tetrachlorophenol	4901513	mg l-1	N	< 0.0002	< 0.0002	< 0.0002	< 0.0002		
	2,3,4,6-Tetrachlorophenol	58902	mg l-1	N	< 0.0002	< 0.0002	< 0.0002	< 0.0002		
	2,3,5,6-Tetrachlorophenol	935955	mg l-1	Ν	< 0.0002	< 0.0002	< 0.0002	< 0.0002		

¹No sampling date was specified, stability times for this analyte may have been exceeded and these results may be compromised. The accreditation for these results remains unaffected.

All tests undertaken between 10/12/2012 and 14/12/2012

* Accreditation status

Column page 1 Report page 3 of 6 LIMS sample ID range Al05103 to Al05259

LABORATORY TEST REPORT



Results of analysis of 4 samples received 10 December 2012

Report Date 14 December 2012

FAO Rob Hooker

Banbury

						217	776	
					AI05256	AI05257	AI05258	AI05259
					HTP02	HTP05	HTP11	HTP35
					8	22	51	57
					Not Provided	Not Provided	Not Provided	Not Provided
					0.50m	0.20m	0.50m	0.60m
					LEACHATE	LEACHATE	LEACHATE	LEACHATE
1900	2,3,4-Trichlorophenol	15950660	mg l-1	N	< 0.0002	<0.0002	< 0.0002	< 0.0002
	2,3,5-Trichlorophenol	933788	mg l-1	Ν	< 0.0002	<0.0002	<0.0002	< 0.0002
	2,3,6-Trichlorophenol	933755	mg l-1	Ν	<0.0002	<0.0002	<0.0002	< 0.0002
	2,4,5-Trichlorophenol	95954	mg l-1	N	<0.0002	<0.0002	<0.0002	< 0.0002
	2,4,6-Trichlorophenol	88062	mg l-1	Ν	< 0.0002	<0.0002	< 0.0002	< 0.0002
	3,4,5-Trichlorophenol	609198	mg l-1	Ν	< 0.0002	< 0.0002	< 0.0002	< 0.0002

LABORATORY TEST REPORT



Results of analysis of 24 samples received 10 December 2012

Report Date 14 December 2012

FAO Rob Hooker

Banbury

Login E	Batch No						217	776		
Chemte	est LIMS ID				AI05105	AI05108	AI05109	AI05112	AI05113	AI05115
Sample	ID				HTP02	HTP03	HTP04	HTP05	HTP06	HTP07
Sample	No				7	12	16	23	28	33
Samplir	ng Date				Not Provided					
Depth					0.20m	0.50m	0.10m	0.50m	0.10m	0.50m
Matrix					SOIL	SOIL	SOIL	SOIL	SOIL	SOIL
SOP↓	Determinand↓	CAS No↓	Units↓	*						
2010	рН			М	6.7	8.3	7.7	8.2	8.0	7.1
2300	Cyanide (free)	57125	mg kg-1	М	<0.50	<0.50	<0.50	<0.50	<0.50	<0.50
2625	Fraction of Organic Carbon			М	0.015	0.0065	0.030	0.0093	0.12	0.023
2120	Boron (hot water soluble)	7440428	mg kg-1	М	0.8	1	1	0.6	1.2	0.5
2490	Chromium (hexavalent)	18540299	mg kg-1	N	<0.5	<0.5	<0.5	<0.5	<0.5	<0.5
2450	Arsenic	7440382	mg kg-1	М	28	84	56	33	48	18
	Beryllium	7440417	mg kg-1	М	<1.00	2.1	1.6	1.1	1.8	<1.00
	Cadmium	7440439	mg kg-1	М	<0.10	0.10	<0.10	<0.10	0.23	<0.10
	Chromium	7440473	mg kg-1	М	47	150	100	75	80	49
	Copper	7440508	mg kg-1	М	18	36	26	18	150	15
	Mercury	7439976	mg kg-1	М	<0.10	<0.10	<0.10	<0.10	0.71	<0.10
	Nickel	7440020	mg kg-1	М	25	64	67	45	53	33
	Lead	7439921	mg kg-1	М	39	230	52	30	480	26
	Selenium	7782492	mg kg-1	М	<0.20	<0.20	<0.20	<0.20	<0.20	<0.20
	Vanadium	7440622	mg kg-1	М	67	230	150	95	120	61
	Zinc	7440666	mg kg-1	М	79	170	190	110	410	85
2800	Naphthalene	91203	mg kg-1	М	< 0.01	0.069	< 0.01	< 0.01	1.8	< 0.01
	Acenaphthylene	208968	mg kg-1	N	< 0.01	0.2	< 0.01	< 0.01	2.8	< 0.01
	Acenaphthene	83329	mg kg-1	М	< 0.01	0.088	< 0.01	< 0.01	0.73	< 0.01
	Fluorene	86737	mg kg-1	М	< 0.01	0.16	< 0.01	< 0.01	2.3	< 0.01
	Phenanthrene	85018	mg kg-1	М	0.088	2.7	0.014	0.049	27	0.018
	Anthracene	120127	mg kg-1	М	< 0.01	0.47	< 0.01	0.013	6	< 0.01
	Fluoranthene	206440	mg kg-1	М	0.17	3.6	0.075	0.2	36	0.047

¹No sampling date was specified, stability times for this analyte may have been exceeded and these results may be compromised. The accreditation for these results remains unaffected.

All tests undertaken between 10/12/2012 and 14/12/2012

* Accreditation status

Column page 1 Report page 5 of 6 LIMS sample ID range Al05103 to Al05259

LABORATORY TEST REPORT



Results of analysis of 24 samples received 10 December 2012

Report Date 14 December 2012

FAO Rob Hooker

Banbury

Logi	n Batch No						217	776		
Chei	mtest LIMS ID				AI05117	AI05118	AI05121	AI05123	AI05125	AI05127
Sam	ple ID				HTP09	HTP10	HTP13	HTP15	HTP17	HTP20
Sam	ple No				41	47	116	101	109	129
Sam	pling Date				Not Provided					
Dep	th				0.20m	0.50m	0.20m	0.25m	0.15m	0.15m
Matr	ix				SOIL	SOIL	SOIL	SOIL	SOIL	SOIL
SOF	P↓ Determinand↓	CAS No↓	Units↓	*						
2010	рН			М	6.9	7.3	7.3	7.1	7.5	7.7
2300	Cyanide (free)	57125	mg kg-1	М	<0.50	<0.50	<0.50	<0.50	<0.50	<0.50
2625	Fraction of Organic Carbon			М	0.0046	0.019	0.018	0.0099	0.017	0.023
2120	Boron (hot water soluble)	7440428	mg kg-1	М	<0.4	0.5	0.7	0.8	0.9	1.4
2490	Chromium (hexavalent)	18540299	mg kg-1	Ν	<0.5	<0.5	<0.5	<0.5	<0.5	< 0.5
2450	Arsenic	7440382	mg kg-1	М	22	18	17	110	66	68
	Beryllium	7440417	mg kg-1	М	<1.00	<1.00	<1.00	4.1	1.6	2.3
	Cadmium	7440439	mg kg-1	Μ	<0.10	<0.10	<0.10	<0.10	<0.10	0.28
	Chromium	7440473	mg kg-1	М	41	42	39	270	78	88
	Copper	7440508	mg kg-1	Μ	12	10	12	21	23	31
	Mercury	7439976	mg kg-1	М	<0.10	<0.10	<0.10	<0.10	<0.10	<0.10
	Nickel	7440020	mg kg-1	Μ	22	22	21	110	62	99
	Lead	7439921	mg kg-1	М	16	24	29	39	46	62
	Selenium	7782492	mg kg-1	Μ	<0.20	<0.20	<0.20	<0.20	<0.20	<0.20
	Vanadium	7440622	mg kg-1	М	58	58	53	410	120	150
	Zinc	7440666	mg kg-1	Μ	59	66	61	190	110	180
2800	Naphthalene	91203	mg kg-1	Μ	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
	Acenaphthylene	208968	mg kg-1	Ν	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
	Acenaphthene	83329	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
	Fluorene	86737	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
	Phenanthrene	85018	mg kg-1	М	< 0.01	< 0.01	0.014	0.039	< 0.01	0.39
	Anthracene	120127	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	0.11
	Fluoranthene	206440	mg kg-1	Μ	< 0.01	0.031	0.088	0.096	0.019	0.81

¹No sampling date was specified, stability times for this analyte may have been exceeded and these results may be compromised. The accreditation for these results remains unaffected.

Column page 2 Report page 5 of 6 LIMS sample ID range Al05103 to Al05259

* Accreditation status

LABORATORY TEST REPORT



Results of analysis of 24 samples received 10 December 2012

Report Date 14 December 2012

FAO Rob Hooker

Banbury

Login Batch No						217	776		
Chemtest LIMS ID				AI05129	AI05131	AI05132	AI05133	AI05135	AI05137
Sample ID				HTP22	HTP24	HTP25	HTP26	HTP29	HTP33
Sample No				137	76	72	68	87	64
Sampling Date				Not Provided					
Depth				0.20m	0.20m	0.15m	0.10m	0.15m	0.25m
Matrix				SOIL	SOIL	SOIL	SOIL	SOIL	SOIL
SOP↓ Determinand↓	CAS No↓	Units↓	*						
2010 pH			М	8.3	7.7	7.7	7.9	7.9	7.6
2300 Cyanide (free)	57125	mg kg-1	М	<0.50	<0.50	<0.50	<0.50	<0.50	< 0.50
2625 Fraction of Organic Carbon			М	0.021	0.057	0.012	0.013	0.023	0.017
2120 Boron (hot water soluble)	7440428	mg kg-1	М	0.7	0.9	0.5	1.1	1.3	1.2
2490 Chromium (hexavalent)	18540299	mg kg-1	Ν	<0.5	< 0.5	<0.5	<0.5	<0.5	<0.5
2450 Arsenic	7440382	mg kg-1	М	86	79	140	120	58	76
Beryllium	7440417	mg kg-1	М	2.5	2.0	4.2	3.7	1.6	2.1
Cadmium	7440439	mg kg-1	М	<0.10	<0.10	<0.10	<0.10	<0.10	<0.10
Chromium	7440473	mg kg-1	М	110	110	270	240	78	100
Copper	7440508	mg kg-1	М	31	31	41	46	28	40
Mercury	7439976	mg kg-1	М	<0.10	<0.10	<0.10	<0.10	<0.10	<0.10
Nickel	7440020	mg kg-1	М	100	92	150	120	69	79
Lead	7439921	mg kg-1	М	55	51	70	120	48	71
Selenium	7782492	mg kg-1	М	<0.20	<0.20	0.22	<0.20	<0.20	<0.20
Vanadium	7440622	mg kg-1	М	150	160	380	320	120	150
Zinc	7440666	mg kg-1	М	140	160	300	260	110	150
2800 Naphthalene	91203	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
Acenaphthylene	208968	mg kg-1	N	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
Acenaphthene	83329	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
Fluorene	86737	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
Phenanthrene	85018	mg kg-1	М	< 0.01	< 0.01	< 0.01	0.043	0.059	0.029
Anthracene	120127	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	0.012	< 0.01
Fluoranthene	206440	mg kg-1	М	0.043	< 0.01	0.028	0.13	0.26	0.076

¹No sampling date was specified, stability times for this analyte may have been exceeded and these results may be compromised. The accreditation for these results remains unaffected.

Column page 3 Report page 5 of 6 LIMS sample ID range Al05103 to Al05259

* Accreditation status

LABORATORY TEST REPORT



Results of analysis of 24 samples received 10 December 2012

FAO Rob Hooker

Banbury

Report Date
14 December 2012

Login Batch No				217	776
Chemtest LIMS ID				AI05140	AI05141
Sample ID				HTP39	HTP40
Sample No			_	90	97
Sampling Date				Not Provided	Not Provided
Depth				0.20m	0.20m
Matrix				SOIL	SOIL
SOP↓ Determinand↓	CAS No↓	Units↓	*		
2010 pH			М	7.6	7.3
2300 Cyanide (free)	57125	mg kg-1	М	<0.50	< 0.50
2625 Fraction of Organic Carbon			М	0.011	0.013
2120 Boron (hot water soluble)	7440428	mg kg-1	М	0.7	0.9
2490 Chromium (hexavalent)	18540299	mg kg-1	Ν	<0.5	<0.5
2450 Arsenic	7440382	mg kg-1	М	52	58
Beryllium	7440417	mg kg-1	М	1.4	1.6
Cadmium	7440439	mg kg-1	М	<0.10	<0.10
Chromium	7440473	mg kg-1	М	78	82
Copper	7440508	mg kg-1	М	23	26
Mercury	7439976	mg kg-1	М	<0.10	<0.10
Nickel	7440020	mg kg-1	М	52	60
Lead	7439921	mg kg-1	М	44	49
Selenium	7782492	mg kg-1	М	<0.20	<0.20
Vanadium	7440622	mg kg-1	М	100	120
Zinc	7440666	mg kg-1	М	100	120
2800 Naphthalene	91203	mg kg-1	М	< 0.01	< 0.01
Acenaphthylene	208968	mg kg-1	Ν	0.073	0.026
Acenaphthene	83329	mg kg-1	М	< 0.01	< 0.01
Fluorene	86737	mg kg-1	М	< 0.01	< 0.01
Phenanthrene	85018	mg kg-1	М	0.28	0.17
Anthracene	120127	mg kg-1	М	0.14	0.064
Fluoranthene	206440	mg kg-1	М	1.7	0.84

¹No sampling date was specified, stability times for this analyte may have been exceeded and these results may be compromised. The accreditation for these results remains unaffected.

Column page 4 Report page 5 of 6 LIMS sample ID range Al05103 to Al05259

* Accreditation status

LABORATORY TEST REPORT



Results of analysis of 24 samples received 10 December 2012

Report Date 14 December 2012

FAO Rob Hooker

Banbury

							217	776		
					AI05105	AI05108	AI05109	AI05112	AI05113	Al05115
					HTP02	HTP03	HTP04	HTP05	HTP06	HTP07
					7	12	16	23	28	33
					Not Provided					
					0.20m	0.50m	0.10m	0.50m	0.10m	0.50m
					SOIL	SOIL	SOIL	SOIL	SOIL	SOIL
2800	Pyrene	129000	mg kg-1	М	0.11	2.9	0.066	0.16	29	0.035
	Benzo[a]anthracene	56553	mg kg-1	М	0.04	1.6	0.031	0.11	17	0.023
	Chrysene	218019	mg kg-1	М	0.027	1.3	0.03	0.061	15	0.013
	Benzo[b]fluoranthene	205992	mg kg-1	М	0.074	1.8	0.047	0.2	20	< 0.01
	Benzo[k]fluoranthene	207089	mg kg-1	Ν	< 0.01	0.64	0.012	0.026	5.7	< 0.01
	Benzo[a]pyrene	50328	mg kg-1	М	0.021	1.3	0.017	0.087	14	< 0.01
	Dibenzo[a,h]anthracene	53703	mg kg-1	Ν	< 0.01	< 0.01	< 0.01	< 0.01	2	< 0.01
	Indeno[1,2,3-cd]pyrene	193395	mg kg-1	М	< 0.01	0.53	< 0.01	0.03	7.2	< 0.01
	Benzo[g,h,i]perylene	191242	mg kg-1	М	< 0.01	0.67	< 0.01	0.033	7.8	< 0.01
	Total (of 16) PAHs		mg kg-1	Ν	0.53	18	0.29	0.97	190	< 0.2
2920	Phenols (total)		mg kg-1	Ν	<0.3	<0.3	<0.3	<0.3	<0.3	<0.3

¹No sampling date was specified, stability times for this analyte may have been exceeded and these results may be compromised. The accreditation for these results remains unaffected.

LABORATORY TEST REPORT



Results of analysis of 24 samples received 10 December 2012

Report Date 14 December 2012

FAO Rob Hooker

* Accreditation status

Banbury

							217	776		
					Al05117	Al05118	AI05121	AI05123	AI05125	AI05127
					HTP09	HTP10	HTP13	HTP15	HTP17	HTP20
					41	47	116	101	109	129
					Not Provided					
					0.20m	0.50m	0.20m	0.25m	0.15m	0.15m
					SOIL	SOIL	SOIL	SOIL	SOIL	SOIL
2800	Pyrene	129000	mg kg-1	М	< 0.01	0.017	0.061	0.037	0.028	0.64
	Benzo[a]anthracene	56553	mg kg-1	М	< 0.01	0.012	0.021	< 0.01	< 0.01	0.36
	Chrysene	218019	mg kg-1	М	< 0.01	0.01	0.012	< 0.01	< 0.01	0.3
	Benzo[b]fluoranthene	205992	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	0.5
	Benzo[k]fluoranthene	207089	mg kg-1	N	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	0.13
	Benzo[a]pyrene	50328	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	0.38
	Dibenzo[a,h]anthracene	53703	mg kg-1	N	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
	Indeno[1,2,3-cd]pyrene	193395	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	0.083
	Benzo[g,h,i]perylene	191242	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	0.12
	Total (of 16) PAHs		mg kg-1	N	< 0.2	< 0.2	< 0.2	< 0.2	< 0.2	3.8
2920	Phenols (total)		mg kg-1	Ν	<0.3	<0.3	<0.3	<0.3	<0.3	<0.3

LABORATORY TEST REPORT



Results of analysis of 24 samples received 10 December 2012

Report Date 14 December 2012

FAO Rob Hooker

* Accreditation status

Banbury

							217	776		
					AI05129	Al05131	AI05132	AI05133	AI05135	Al05137
					HTP22	HTP24	HTP25	HTP26	HTP29	HTP33
					137	76	72	68	87	64
					Not Provided					
					0.20m	0.20m	0.15m	0.10m	0.15m	0.25m
					SOIL	SOIL	SOIL	SOIL	SOIL	SOIL
2800	Pyrene	129000	mg kg-1	М	0.03	< 0.01	0.024	0.1	0.21	0.056
	Benzo[a]anthracene	56553	mg kg-1	М	< 0.01	< 0.01	< 0.01	0.047	0.099	0.023
	Chrysene	218019	mg kg-1	М	< 0.01	< 0.01	< 0.01	0.029	0.08	0.024
	Benzo[b]fluoranthene	205992	mg kg-1	М	< 0.01	< 0.01	0.01	0.046	0.12	0.021
	Benzo[k]fluoranthene	207089	mg kg-1	N	< 0.01	< 0.01	< 0.01	< 0.01	0.011	< 0.01
	Benzo[a]pyrene	50328	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	0.07	0.018
	Dibenzo[a,h]anthracene	53703	mg kg-1	N	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
	Indeno[1,2,3-cd]pyrene	193395	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
	Benzo[g,h,i]perylene	191242	mg kg-1	М	< 0.01	< 0.01	< 0.01	< 0.01	0.013	< 0.01
	Total (of 16) PAHs		mg kg-1	N	< 0.2	< 0.2	< 0.2	0.4	0.93	0.25
2920	Phenols (total)		mg kg-1	Ν	<0.3	<0.3	<0.3	<0.3	<0.3	<0.3

LABORATORY TEST REPORT

Results of analysis of 24 samples received 10 December 2012



Report Date 14 December 2012

FAO Rob Hooker

* Accreditation status

Banbury

					217	776
					AI05140	AI05141
						HTP40
					90	97
					Not Provided	Not Provided
					0.20m	0.20m
					SOIL	SOIL
2800	Pyrene	129000	mg kg-1	М	1.4	0.65
	Benzo[a]anthracene	56553	mg kg-1	М	0.95	0.48
	Chrysene	218019	mg kg-1	М	0.78	0.32
	Benzo[b]fluoranthene	205992	mg kg-1	М	1.2	0.62
	Benzo[k]fluoranthene	207089	mg kg-1	N	0.35	0.099
	Benzo[a]pyrene	50328	mg kg-1	М	0.69	0.33
	Dibenzo[a,h]anthracene	53703	mg kg-1	N	< 0.01	< 0.01
	Indeno[1,2,3-cd]pyrene	193395	mg kg-1	М	0.22	0.069
	Benzo[g,h,i]perylene	191242	mg kg-1	М	0.21	0.076
	Total (of 16) PAHs		mg kg-1	N	8	3.7
2920	Phenols (total)		mg kg-1	Ν	<0.3	< 0.3



Depot Road Newmarket CB8 0AL Tel: 01638 606070

Hydrock Consultants Over Court Barns Over Lane Almondsbury, Bristol BS32 4DF

FAO Richard Heath 15 March 2013

Dear Richard Heath

Test Report Number	225356
Your Project Reference	C12702 - Bankside, Banbury

Please find enclosed the results of analysis for the samples received 8 March 2013.

All soil samples will be retained for a period of one month and all water samples will be retained for 7 days following the date of the test report. Should you require an extended retention period then please detail your requirements in an email to customerservices@chemtest.co.uk. Please be aware that charges may be applicable for extended sample storage.

If you require any further assistance, please do not hesitate to contact the Customer Services team.

Yours sincerely



Darrell Hall, Director



Notes to accompany report:

- The sign < means 'less than'
- Tests marked 'U' hold UKAS accreditation
- Tests marked 'M' hold MCertS (and UKAS) accreditation
- Tests marked 'N' do not currently hold UKAS accreditation
- Tests marked 'S' were subcontracted to an approved laboratory
- n/e means 'not evaluated'
- i/s means 'insufficient sample'
- u/s means 'unsuitable sample'
- Comments or interpretations are beyond the scope of UKAS accreditation
 - The results relate only to the items tested
- All results are expressed on a dry weight basis
- The following tests were analysed on samples as received and the results subsequently corrected to a dry weight basis TPH, BTEX, VOCs, SVOCs, PCBs, phenols
 - For all other tests the samples were dried at < 37°C prior to analysis
 - Uncertainties of measurement for the determinands tested are available upon request
 - None of the test results included in this report have been recovery corrected

Test Report 225356 Cover Sheet



Depot Road Newmarket CB8 0AL Tel: 01638 606070

Hydrock Consultants Over Court Barns Over Lane Almondsbury, Bristol BS32 4DF

FAO Richard Heath 02 April 2013

Dear Richard Heath

Test Report Number225356Your Project ReferenceC12702

C12702 - Bankside, Banbury

Please find enclosed the results of analysis for the samples received 8 March 2013.

If you require any further assistance, please do not hesitate to contact the Customer Services team.

Yours sincerely

AD

Keith Jones, Technical Manager

Notes to accompany report:

- The in-house procedure is employed to identify materials and fibres in soils
- The sample is examined by stereo-binocular and polarised light microscopy
- Sample size is reduced by coning and quartering to obtain a representative sub-sample if necessary
- The bulk identification is in accordance with the requirements of the analyst guide (HSG 248)
- Samples associated with asbestos are retained for six months
- The results relate only to the items tested as supplied by the client
 - Comments or interpretations are beyond the scope of UKAS accreditation



Test Report 225356 Cover Sheet

Newmarket • Coventry • Dublin Registered in England & Wales - Registration Number 6511736 - Registered Office: 11 Depot Road Newmarket Suffolk CB8 0AL

LABORATORY TEST REPORT Asbestos in Soils

Chemtest 6

FAO Richard Heath

Results of analysis of 5 samples received 8 March 2013 C12702 - Bankside, Banbury

Report Date 02 April 2013

Login Batch No: 225356

Qualitative Results

				SO	P 2190
				АСМ Туре	Asbestos Identification
Chemtest ID	Sample ID	Sample Desc	Depth (m)		
AI39854	HTP41		0.1	-	No Asbestos Detected
AI39856	HTP43		0.2	-	No Asbestos Detected
AI39857	HTP44		0.7	-	No Asbestos Detected
AI39858	HTP45		0.4	-	No Asbestos Detected
AI39860	HTP46		0.7	-	No Asbestos Detected

The detection limit for this method is 0.001%

Signed

Albert Vella Senior Environmental Surveyor

LABORATORY TEST REPORT



Results of analysis of 5 samples received 8 March 2013

Report Date 14 March 2013

FAO Richard Heath

C12702 - Bankside, Banbury

Login E	Batch No						225356		
Chemte	est LIMS ID				AI39854	AI39856	AI39857	AI39858	AI39860
Sample	ID				HTP41	HTP43	HTP44	HTP45	HTP46
Sample	No								
Sampli	ng Date				Not Provided				
Depth					0.1m	0.2m	0.7m	0.4m	0.7m
Matrix					SOIL	SOIL	SOIL	SOIL	SOIL
SOP↓	Determinand↓	CAS No↓ U	nits↓ *						
2010	рН			М	7.7	7.5	7.6	6.9	7.9
2300	Cyanide (free)	57125	mg kg-1	М	<0.5	<0.5	<0.5	<0.5	<0.5
2625	Fraction of Organic Carbon			М	0.023	0.018	0.012	0.022	0.0053
2120	Boron (hot water soluble)	7440428	mg kg-1	М	1.0	1.0	0.7	1.0	1.2
2490	Chromium (hexavalent)	18540299	mg kg-1	Ν	<0.5	<0.5	<0.5	<0.5	<0.5
2450	Arsenic	7440382	mg kg-1	М	190	130	230	120	170
	Beryllium	7440417	mg kg-1	М	5.4	3.6	5.6	3.3	4.8
	Cadmium	7440439	mg kg-1	М	0.36	0.37	0.27	0.40	0.23
	Chromium	7440473	mg kg-1	М	480	210	540	190	590
	Copper	7440508	mg kg-1	М	40	49	35	43	25
	Mercury	7439976	mg kg-1	М	0.19	0.19	0.13	0.17	<0.10
	Nickel	7440020	mg kg-1	М	170	160	210	130	180
	Lead	7439921	mg kg-1	М	81	90	84	81	33
	Selenium	7782492	mg kg-1	М	<0.20	<0.20	<0.20	<0.20	<0.20
	Vanadium	7440622	mg kg-1	М	670	330	710	290	740
	Zinc	7440666	mg kg-1	М	340	290	380	220	970
2800	Naphthalene	91203	mg kg-1	М	0.8	1.5	0.95	1.2	2
	Acenaphthylene	208968	mg kg-1	N	< 0.01	0.063	0.03	0.028	< 0.01
	Acenaphthene	83329	mg kg-1	М	< 0.01	0.1	0.044	0.054	0.091
	Fluorene	86737	mg kg-1	М	< 0.01	0.08	0.038	0.053	0.11
	Phenanthrene	85018	mg kg-1	М	0.22	0.17	0.1	0.08	0.24
	Anthracene	120127	mg kg-1	М	0.03	0.02	< 0.01	< 0.01	0.031
	Fluoranthene	206440	mg kg-1	М	0.24	0.17	0.091	0.071	0.096
	Pyrene	129000	mg kg-1	М	0.17	0.12	0.072	0.045	0.063

LABORATORY TEST REPORT



Results of analysis of 5 samples received 8 March 2013

Report Date 14 March 2013

FAO Richard Heath

C12702 - Bankside, Banbury

							225356		
					AI39854	AI39856	AI39857	AI39858	AI39860
					HTP41	HTP43	HTP44	HTP45	HTP46
					Not Provided				
					0.1m	0.2m	0.7m	0.4m	0.7m
					SOIL	SOIL	SOIL	SOIL	SOIL
2800	Benzo[a]anthracene	56553	mg kg-1	М	0.11	0.04	< 0.01	< 0.01	< 0.01
	Chrysene	218019	mg kg-1	M	0.11	0.054	< 0.01	< 0.01	< 0.01
	Benzo[b]fluoranthene	205992	mg kg-1	М	0.18	0.056	< 0.01	< 0.01	< 0.01
	Benzo[k]fluoranthene	207089	mg kg-1	N	0.014	0.023	< 0.01	< 0.01	< 0.01
	Benzo[a]pyrene	50328	mg kg-1	М	0.055	0.023	< 0.01	< 0.01	< 0.01
	Dibenzo[a,h]anthracene	53703	mg kg-1	Ν	< 0.01	< 0.01	< 0.01	< 0.01	< 0.01
	Indeno[1,2,3-cd]pyrene	193395	mg kg-1	М	0.025	< 0.01	< 0.01	< 0.01	< 0.01
	Benzo[g,h,i]perylene	191242	mg kg-1	М	0.032	< 0.01	< 0.01	< 0.01	< 0.01
	Total (of 16) PAHs		mg kg-1	Ν	2	2.4	1.3	1.5	2.6
2920	Phenols (total)		mg kg-1	N	<0.3	<0.3	<0.3	<0.3	<0.3

All tests undertaken between 08/03/2013 and 14/03/2013

* Accreditation status

Column page 1 Report page 2 of 2 LIMS sample ID range Al39854 to Al39860

Chemical of Potential Concern Lab. RL Arsenic 2 Beryllium 1 Boron 0.4 Cadmium 0.1 Chromium (III) 5	L N Sam		Min.	Max.	No.			Location & Depth	HTP02 0.20	HTP03 0.50	HTP04 0.10	HTP05	HTP06	TP1	TP1	TP2	TP2	TP2	TP3	TP3	TP3	TP4	TP4	TP7	TP7	TP10	TP10	TP11	TP12	TP12
Concern Lab. RL Arsenic 2 Beryllium 1 Boron 0.4 Cadmium 0.1 Chromium (III) 5	Sam								0.20	0.50	0.40	0.50																		
Beryllium 1 Boron 0.4 Cadmium 0.1 Chromium (III) 5	2			Value	Samples > or = GAC	GAC	US ₉₅	Result of Significance Test		0.50	0.10	0.50	0.10	0.4	0.90m	0.2	0.5	1.5	0.1	0.5	1.8	0.6	1.5	0.2	0.7	0.3	0.9	0.3	0.2	0.4
Boron 0.4 Cadmium 0.1 Chromium (III) 5		28	12	150	19	32	79.77682	FURTHER ASSESSMENT REQUIRED	28	84	56	33	48	150	76		26	22	42	40	78	27	22	72	72		45	49	69	66
Cadmium 0.1 Chromium (III) 5		7	0.5	2.1	0	51	2.260837	POTENTIALLY SUITABLE FOR USE	1	2.1	1.6	1.1	1.8	0.5	0.5															
Chromium (III) 5	2	28	0.6	3.8	0	290	2.162511	POTENTIALLY SUITABLE FOR USE	0.8	1	1	0.6	1.2	1.6	1.6		2.3	0.86	2.8	1.4	2.2	1.7	0.68	2.4	3.8		1	2.8	1.2	1.2
			0.1	0.71	0	11	0.576684	POTENTIALLY SUITABLE FOR USE	0.1	0.1	0.1	0.1	0.23	0.61	0.5		0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.52		0.5	0.71	0.5	0.5
		28 :	31.9	410	0	630	159.8763	POTENTIALLY SUITABLE FOR USE	46.5	149.5	99.5	74.5	79.5	410	159.9		38.89	31.9	74.9	63	129.9	62.88	37.67	80	109.85		39	72.85	66	129.9
Chromium (VI) 0.5			0.1	0.5	0	4.3	0.41444	POTENTIALLY SUITABLE FOR USE	0.5	0.5	0.5	0.5	0.5	1	0.1		0.11	0.1	<u>0</u> .1		0.1	0.12	0.33		0.15			0.15	1	0.1
Copper 5			7.3	230	0	2300	87.18576	POTENTIALLY SUITABLE FOR USE	18	36	26	18	150	7.3	33		23	13	180	23	16	21	19	25	110		24	230	25	17
Lead 5			17	1500	2	450	381.0785	POTENTIALLY SUITABLE FOR USE	39	230	52	30	480	40	75		62	17	410	24	33	51	20	64	360		29	1500	40	40
Mercury, inorganic 0.1			0.1	2.6	0	170	1.029042	POTENTIALLY SUITABLE FOR USE	0.1	0.1	0.1	0.1	0.71	0.27	0.92		0.65	0.24	2.1	0.2	0.39	0.21	0.2	0.27	2.6		0.2	2	0.2	0.25
Nickel 5		28	23	150	1	130	86.43294	POTENTIALLY SUITABLE FOR USE	25	64	67	45	53	150	80		23	28	52	80	59	31	33	81	64		70	77	110	65
Selenium 0.2	2	28	0.2	0.76	0	350	0.431448	POTENTIALLY SUITABLE FOR USE	0.2	0.2	0.2	0.2	0.2	0.3	0.38		0.3	0.3	0.49	0.3	0.33	0.32	0.3	0.3	0.76		0.3	0.59	0.43	0.3
Vanadium 5			0.5	230	4		230.1368	FURTHER ASSESSMENT REQUIRED	67	230	150	95	120	0.5	0.5												1			
Zinc 10			56	410	0		229.6131	POTENTIALLY SUITABLE FOR USE	79	170	190	110	410	260	180		130	64	300	59	160	98	64	130	260		64	320	120	150
Cyanide (free) 0.5	2	28	0.5	5	0	750	4.775601	POTENTIALLY SUITABLE FOR USE	0.5	0.5	0.5	0.5	0.5	2	5		5	5	5	2	5	5	5	2	5		2	5	2	5
Phenol (total) 0.3			0.3	4.8	0	290	1.948196	POTENTIALLY SUITABLE FOR USE	0.3	0.3	0.3	0.3	0.3	0.5	4.3		0.5	0.5	3.7	0.5	0.5	0.5	0.5	0.5	4.8		0.5	2.1	0.5	0.5
Acenaphthene 0.01	1	19 1	0.01	11	0	480	3.549764	POTENTIALLY SUITABLE FOR USE	0.01	0.088	0.01	0.01	0.73		0.5		0.5	0.5	0.5		0.5	0.5	0.5		11			2.7		0.5
Acenaphthylene 0.01	1	19 0	0.01	5.4	0	400	2.106977	POTENTIALLY SUITABLE FOR USE	0.01	0.2	0.01	0.01	2.8	1	0.5	[0.5	0.5	5.4		0.5	0.5	0.5		1.5		1	0.51	1	0.5
Anthracene 0.01	1		0.01	19	0	4900	8.456297	POTENTIALLY SUITABLE FOR USE	0.01	0.47	0.01	0.013	6	1	0.92		0.85	0.5	17		0.5	0.5	0.5		19		1	5.4	1	0.5
Benz(a)anthracene 0.01			0.031	36	4	4.7	16.01503	FURTHER ASSESSMENT REQUIRED	0.04	1.6	0.031	0.11	17		2		2.3	0.5	29		0.5	0.74	0.5		36			11		0.5
Benzo(a)pyrene 0.01			0.017	56	9	0.94	20.05253	FURTHER ASSESSMENT REQUIRED	0.021	1.3	0.017	0.087	14	1	2.3		2.8	0.5	25		0.5	1.1	0.5		56			13	1	0.5
Benzo(b)fluoranthene 0.01	1	19 0	0.047	40	4	6.5	16.18368	FURTHER ASSESSMENT REQUIRED	0.074	1.8	0.047	0.2	20		1.8		2.1	0.5	23		0.5	0.66	0.5		40			9.6		0.5
Benzo(ghi)perylene 0.01			0.01	45	0	46	15.13916	POTENTIALLY SUITABLE FOR USE	0.01	0.67	0.01	0.033	7.8	1	1.7		2.2	0.5	15		0.5	1	0.5		45			9.1	1	0.5
Benzo(k)fluoranthene 0.01	1		0.01	26	2	9.6	9.280907	POTENTIALLY SUITABLE FOR USE	0.01	0.64	0.012	0.026	5.7	1	0.93		1.1	0.5	11		0.5	0.5	0.5		26			6.8		0.5
Chrysene 0.01		19 0	0.027	40	4	8	18.44244	FURTHER ASSESSMENT REQUIRED	0.027	1.3	0.03	0.061	15	1	2.4		2.7	0.5	38		0.5	0.88	0.5		40		1	12	1	0.5
Dibenz(a,h)anthracene 0.01			0.01	4.5	4	0.86	1.832813	FURTHER ASSESSMENT REQUIRED	0.01	0.01	0.01	0.01	2		0.5		0.5	0.5	1.8		0.5	0.5	0.5		4.5			1.1		0.5
Fluoranthene 0.01	1	19 0	0.075	99	0	460	44.88063	POTENTIALLY SUITABLE FOR USE	0.17	3.6	0.075	0.2	36	1	4.9		6.2	0.5	91		0.5	2.2	0.5		99			31	1	0.5
Fluorene 0.01		19 (0.01	18	0	380	6.306193	POTENTIALLY SUITABLE FOR USE	0.01	0.16	0.01	0.01	2.3	1	0.5		0.5	0.5	18		0.5	0.5	0.5		9.1		1	1.3	1	0.5
Indeno(1,2,3,cd)pyrene 0.01	1	19 (0.01	43	4	3.9	14.60898	FURTHER ASSESSMENT REQUIRED	0.01	0.53	0.01	0.03	7.2		1.4		1.8	0.5	15		0.5	0.91	0.5		43			10		0.5
Naphthalene 0.01			0.01	42	3	3.7	13.63966	FURTHER ASSESSMENT REQUIRED	0.01	0.069	0.01	0.01	1.8	1	1.3		0.91	0.5	42		0.5	0.5	0.5		15		1	5.2	1	0.5
Phenanthrene 0.01			0.014	84	0	200	34.56433	POTENTIALLY SUITABLE FOR USE	0.088	2.7	0.014	0.049	27	1	3		2.6	0.5	84		0.5	1.3	0.5		64			19		0.5
Pyrene 0.01		19 0	0.066	95	0	1000	38.88174	POTENTIALLY SUITABLE FOR USE	0.11	2.9	0.066	0.16	29	1	4.1		4.7	0.5	69		0.5	1.8	0.5		95			27	1	0.5
Mean																														
FOC (dimensionless) 0.018									0.015	0.0065	0.03	0.0093	0.12	0.004244		0.02093				0.004244				0.022674		0.023256			0.006395	
SOM (calculated) 3.17%			1						2.59%	1.12%	5.17%	1.60%	20.69%	0.73%		3.61%	1			0.73%				3.91%		4.01%	0.73%		1.10%	
pH (su) 7.9	1	1							6.7	8.3	7.7	8.2	8	7.9	8.2	7.7	8.6	7.7	8.2	8.4	7.7	8	7.6	7.7	8.1	7.1	8.3	8.1	8.1	8.1

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Client: Bovis Barratt and Taylor Wimpey Site: Land at Bankside, Banbury Job no: C12702

Values in red are equal to, or greater than, the generic assessment criterion (GAC).

MG denotes Made Ground

NAT denotes natural ground

Hydrock

							Soil Type									
l values i	n mg/kg unle	ss otherwis	e stated				Location & Depth	TP13	TP13	TP13	TP14	TP15	TP16	TP16	TP23	TP24
.ab. RL	No. Samples	Min. Value	Max. Value	No. Samples > or = GAC	GAC	US ₉₅	Result of Significance Test	0.3	0.8	1	0.4	2.7	0.4	0.7	0.4	0.4
2	28	12	150	19	32	79.77682	FURTHER ASSESSMENT REQUIRED	82	95		12	16	19	80	21	79
1							POTENTIALLY SUITABLE FOR USE								1	
0.4		0.6	3.8	0	290	2.162511	POTENTIALLY SUITABLE FOR USE	2.2	2.1				0.89	1.2	0.88	0.81
0.1		0.1	0.71	0	11	0.576684	POTENTIALLY SUITABLE FOR USE	0.5							0.5	0.5
		31.9	410	0			POTENTIALLY SUITABLE FOR USE	94					53		40	89
		0.1	0.5	0			POTENTIALLY SUITABLE FOR USE								1	
							POTENTIALLY SUITABLE FOR USE									10
5		17		2			POTENTIALLY SUITABLE FOR USE									25
0.1		0.1		0			POTENTIALLY SUITABLE FOR USE									0.2
5	28	23	150	1			POTENTIALLY SUITABLE FOR USE	91	73		40	29		96	24	86
0.2						0.431448	POTENTIALLY SUITABLE FOR USE	0.42	0.33		0.3	0.34	0.3	0.3	0.3	0.3
5			230	4	74	230.1368	FURTHER ASSESSMENT REQUIRED									
10			410	0	3700	229.6131	POTENTIALLY SUITABLE FOR USE	170	180		120	56	110	150	62	220
0.5	28	0.5	5	0	750	4.775601	POTENTIALLY SUITABLE FOR USE	2	5		5	5	2	5	2	2
0.3	28	0.3	4.8	0	290	1.948196	POTENTIALLY SUITABLE FOR USE	0.5	0.58		0.5	0.76	0.5	0.5	0.5	0.5
0.01	19	0.01	11	0	480	3.549764	POTENTIALLY SUITABLE FOR USE		0.5		0.5	0.5		0.5		
0.01	19	0.01	5.4	0	400	2.106977	POTENTIALLY SUITABLE FOR USE		0.5		0.5	0.5		0.5	1	
0.01	19	0.01	19	0	4900	8.456297	POTENTIALLY SUITABLE FOR USE		0.58		0.5	0.5		0.5	1	
0.01	19	0.031	36	4	4.7	16.01503	FURTHER ASSESSMENT REQUIRED		1.3		0.5	0.5		0.5		
0.01	19	0.017	56	9	0.94	20.05253	FURTHER ASSESSMENT REQUIRED		1.4		0.5	0.5		0.5	1	
0.01	19	0.047	40	4	6.5	16.18368	FURTHER ASSESSMENT REQUIRED		0.97		0.5	0.5		0.5		
0.01	19	0.01	45	0	46	15.13916	POTENTIALLY SUITABLE FOR USE		0.91		0.5	0.5		0.5	1	
0.01	19	0.01	26	2	9.6	9.280907	POTENTIALLY SUITABLE FOR USE		0.77		0.5	0.5		0.5		
0.01	19	0.027	40	4	8	18.44244	FURTHER ASSESSMENT REQUIRED		1.4		0.5	0.5		0.5		
0.01	19	0.01	4.5	4	0.86	1.832813	FURTHER ASSESSMENT REQUIRED		0.5		0.5	0.5		0.5		
0.01	19	0.075	99	0	460	44.88063	POTENTIALLY SUITABLE FOR USE		3.6		0.5	0.5		0.5	1	
0.01	19	0.01	18	0	380	6.306193	POTENTIALLY SUITABLE FOR USE		0.5		0.5	0.5		0.5	1	
0.01	19	0.01	43	4	3.9	14.60898	FURTHER ASSESSMENT REQUIRED		0.98		0.5	0.5		0.5		
0.01	19	0.01	42	3	3.7	13.63966	FURTHER ASSESSMENT REQUIRED		1.1		0.5	0.5		0.5		
0.01	19	0.014	84	0	200	34.56433	POTENTIALLY SUITABLE FOR USE		2.4		0.5	0.5		0.5		
0.01	19	0.066	95	0	1000	38.88174	POTENTIALLY SUITABLE FOR USE		3.1		0.5	0.5		0.5	1	1
Mean										1						
0.018			1	1				0.027907		0.003895			0.003488		0.00814	0.001919
3.17%			1	1		1		4.81%		0.67%			0.60%		1.40%	0.33%
7.9			1	1				8	8.3	8.3	7.3	7.9	7.9	8.2	7.8	8
	2 1 0.4 0.5 5 5 5 5 5 5 5 5 5 5 5 5 5	bb. RL Samples 2 28 1 7 0.4 28 0.1 28 5 28 0.5 109 6 28 0.5 28 0.1 28 5 28 0.2 28 5 7 10 28 0.5 28 0.5 28 0.5 28 0.6 19 0.01 19 0.01 19 0.01 19 0.01 19 0.01 19 0.01 19 0.01 19 0.01 19 0.01 19 0.01 19 0.01 19 0.01 19 0.01 19 0.01 19 0.01 19 0.01 19	ab. RL Samples Value 2 28 12 1 7 0.5 0.4 28 0.6 0.1 28 0.1 5 28 31.9 0.5 10 0.1 5 28 7.3 5 28 17 6 28 7.3 5 28 23 0.2 28 0.1 5 28 23 0.2 28 0.2 5 7 0.5 0.5 28 0.5 0.5 28 0.5 0.5 28 0.5 0.5 28 0.5 0.5 28 0.5 0.6 9 0.01 0.01 19 0.01 0.01 19 0.01 0.01 19 0.01 0.01 19 0.01 0.0	ab, RL Samples Value Value 2 28 12 150 1 7 0.5 2.1 0.4 28 0.6 3.8 0.1 28 0.1 0.7 5 28 31.9 400 0.5 19 0.1 0.5 6 28 7.3 230 0.1 28 0.1 0.5 5 28 17 1500 5 28 17 1500 0.1 28 0.1 2.6 5 28 23 150 0.2 28 0.2 0.76 5 7 0.5 230 0.5 28 0.5 5 0.3 28 0.3 4.8 0.01 19 0.01 19 0.01 19 0.01 240 0.01 19 0.01 40	$\begin{array}{c c c c c c c c c c c c c c c c c c c $	$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	$ \begin{array}{c c c c c c c c c c c c c c c c c c c $	No. Samples Num. Value Max. Value Samples > or = GAC GAC USs Result of Significance Test 2 28 12 150 19 32 79.77682 PURTHER ASSESSMENT REQUERCING 1 70.5 2.1 0 51 2.208037 PORTHALLY SUTABLE FOR USE 0.4 28 0.6 3.8 0 290 2.162511 PORTHALLY SUTABLE FOR USE 0.4 28 0.61 0.71 0 11 0.576684 PORTHALLY SUTABLE FOR USE 5 28 7.3 230 0 2430 0.41444 PORTHALLY SUTABLE FOR USE 5 28 7.3 230 0 2450 381.0785 PORTHALLY SUTABLE FOR USE 5 28 17 1500 2 450 381.0785 PORTHALLY SUTABLE FOR USE 5 28 0.2 0.4 7.4 20.31668 PORTHALLY SUTABLE FOR USE 5 28 0.2 0.76 0 350 4.774 20.3166 PORTHALLY SUTABLE F	No. Samples No. Value Result of Significance Test 2 12 12 150 19 32 79.77682 Further Assessment recover 0 82 0.4 28 0.6 3.8 0 290 2.162511 POTENTIALY SUMALE FOR USE 2.2 5 28 0.1 0.7/1 0 11 0.5630 199.071 0.771.05 2.1 5 28 7.3 230 0 2.300 87.188763 POTENTIALY SUMALE FOR USE 74 0.1 28 0.1 2.6 0 170 1.030424 POTENTIALY SUMALE FOR USE 74 0.1 28 56 410 0 3700 229.17 1.01 2.2 2.3 1.02 2.3 2.3 1.02 2.3 2.4 74 74.73 2.3 <td>bb. RL Samples No. Value Mo. Value No. Value No. Value</td> <td>bb. RL No. Samples No. Value No. Value No. Value No. Value No. Value No. Value No. Soft 2 CAC US_B Result of Significance Test 2 28 12 150 19 32 79.77682 FURMER ASSESSMENT REQUEED 82 95 1 7 0.5 2.1 0 51 2.260837 POTENUALLY SUTABLE FOR USE 2.1 0 0.1 28 0.1 0.6 3.8 0 290 2.162511 POTENUALLY SUTABLE FOR USE 0.5 0.5 0.5 10.5 0.5</td> <td>bb. RL No. Samples No. Value No. Value No. Value No. Value No. Value No. Value Samples > or = GAC GAC US_B Result of Significance Test Image: Comparison of Compariso</td> <td>bb. RL No. Samples Win. Value Max. Value No. Value Solution GAC US_B Result of Significance Test Image: Constraint of Significance Test Image: Constraint of Significance Test 2 28 12 150 19 32 79.77682 FURTHER ASSESSMENT REQUERED 82 95 12 16 1 7 0.5 2.1 0 51 2.260837 POTENNALLY SUTMALE FOR USE 2.2 2.1 0.94 1.3 0.1 28 0.6 3.8 0 290 2.162511 POTENNALLY SUTMALE FOR USE 0.5</td> <td>bb. RL No. Samples Waine No. Value No. Value No. Value Co. Value GAC US₃₀ Result of Significance Test Image: Constraint of the constraint of</td> <td>bb. RL Samples No. Value Max. Value No. Value No. Value</td> <td>bb. Rl No. Samples Winu Watu No. Yalue Solver GAC USs Result of Significance Test Image: Constraint of the solver sector sector of the solver sector of the solver sector of the solver sector of the solver sector sector sector of the solver sector se</td>	bb. RL Samples No. Value Mo. Value No. Value No. Value	bb. RL No. Samples No. Value No. Value No. Value No. Value No. Value No. Value No. Soft 2 CAC US _B Result of Significance Test 2 28 12 150 19 32 79.77682 FURMER ASSESSMENT REQUEED 82 95 1 7 0.5 2.1 0 51 2.260837 POTENUALLY SUTABLE FOR USE 2.1 0 0.1 28 0.1 0.6 3.8 0 290 2.162511 POTENUALLY SUTABLE FOR USE 0.5 0.5 0.5 10.5 0.5	bb. RL No. Samples No. Value No. Value No. Value No. Value No. Value No. Value Samples > or = GAC GAC US _B Result of Significance Test Image: Comparison of Compariso	bb. RL No. Samples Win. Value Max. Value No. Value Solution GAC US _B Result of Significance Test Image: Constraint of Significance Test Image: Constraint of Significance Test 2 28 12 150 19 32 79.77682 FURTHER ASSESSMENT REQUERED 82 95 12 16 1 7 0.5 2.1 0 51 2.260837 POTENNALLY SUTMALE FOR USE 2.2 2.1 0.94 1.3 0.1 28 0.6 3.8 0 290 2.162511 POTENNALLY SUTMALE FOR USE 0.5	bb. RL No. Samples Waine No. Value No. Value No. Value Co. Value GAC US ₃₀ Result of Significance Test Image: Constraint of the constraint of	bb. RL Samples No. Value Max. Value No. Value No. Value	bb. Rl No. Samples Winu Watu No. Yalue Solver GAC USs Result of Significance Test Image: Constraint of the solver sector sector of the solver sector of the solver sector of the solver sector of the solver sector sector sector of the solver sector se

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Assessment of Chemicals of Potential Concern to Plant Life



								Soil Type	MG	NAT	MG	MG	MG										
	All values ir	n mg/kg unles	s otherwise	stated				Location & Depth	HTP02	HTP03	HTP04	HTP05	HTP06	TP1	TP1	TP2	TP2	TP2	TP3	TP3	TP3	TP4	TP4
Chemical of Potential Concern	Lab. RL	No. Samples	Min. Value	Max. Value	No. Samples > or = GAC	GAC	US ₉₅	Result of Significance Test	0.20	0.50	0.10	0.50	0.10	0.4	0.90m	0.2	0.5	1.5	0.1	0.5	1.8	0.6	1.5
Arsenic	2	28	12	150	0	250	79.77682	POTENTIALLY SUITABLE FOR USE	28	84	56	33	48	150	76		26	22	42	40	78	27	22
Boron	0.4	28	0.6	3.8	1	3	2.162511	POTENTIALLY SUITABLE FOR USE	0.8	1	1	0.6	1.2	1.6	1.6		2.3	0.86	2.8	1.4	2.2	1.7	0.68
Chromium (III)	5	28	31.9	410	1	400	159.8763	POTENTIALLY SUITABLE FOR USE	46.5	149.5	99.5	74.5	79.5	410	159.9		38.89	31.9	74.9	63	129.9	62.88	37.67
Chromium (VI)	0.5	19	0.1	0.5	0	25	0.41444	POTENTIALLY SUITABLE FOR USE	0.5	0.5	0.5	0.5	0.5		0.1		0.11	0.1	0.1		0.1	0.12	0.33
Copper	5	28	7.3	230	1	200	87.18576	POTENTIALLY SUITABLE FOR USE	18	36	26	18	150	7.3	33		23	13	180	23	16	21	19
Nickel	5	28	23	150	2	110	86.43294	POTENTIALLY SUITABLE FOR USE	25	64	67	45	53	150	80		23	28	52	80	59	31	33
Zinc	10	28	56	410	3	300	229.6131	POTENTIALLY SUITABLE FOR USE	79	170	190	110	410	260	180		130	64	300	59	160	98	64
	Mean																						
pH (su)	7.9								6.7	8.3	7.7	8.2	8	7.9	8.2	7.7	8.6	7.7	8.2	8.4	7.7	8	7.6

Risk parameter: Plant life pH >7

Data set: Tip Area (COMBINED) Client: Bovis Barratt and Taylor Wimpey Site: Land at Bankside, Banbury Job no: C12702

Legend: Values in blue are at or below the laboratory reporting limit (where a single value is indicated) and are

considered as being at the detection limit for the purposes of statistical analysis, as a conservative estimate.

Values in red are equal to, or greater than, the generic assessment criterion (GAC).

MG denotes Made Ground

NAT denotes natural ground

Assessment of Chemicals of Potential Concern to Plant Life

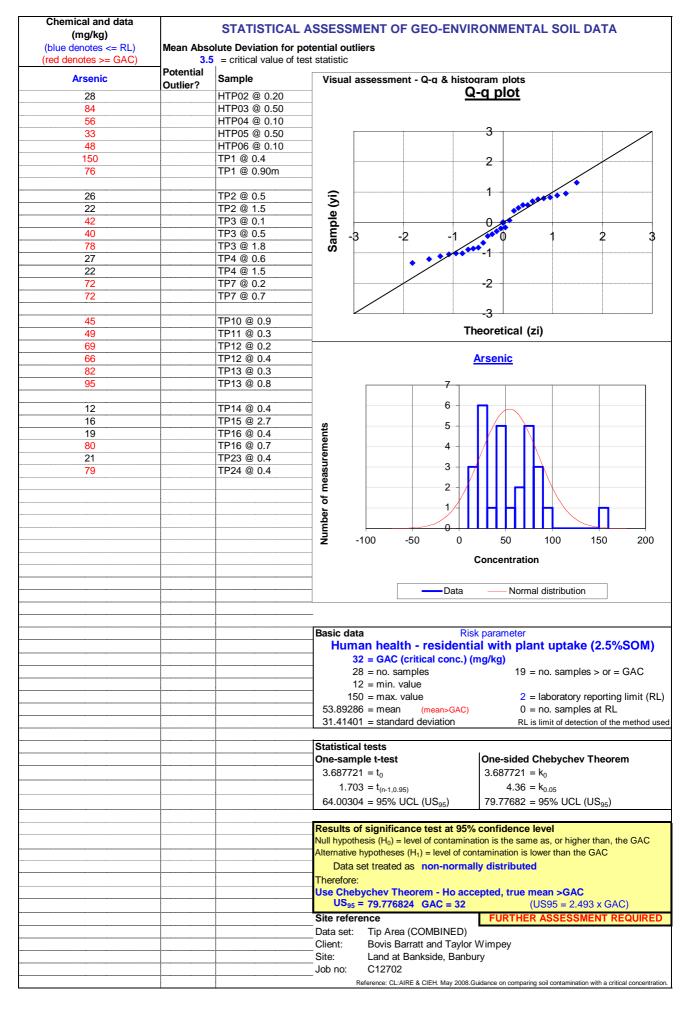


								Soil Type																1
	All values in	n mg/kg unles	s otherwise	stated				Location & Depth	TP7	TP7	TP10	TP10	TP11	TP12	TP12	TP13	TP13	TP13	TP14	TP15	TP16	TP16	TP23	TP24
Chemical of Potential Concern	Lab. RL	No. Samples	Min. Value	Max. Value	No. Samples > or = GAC	GAC	US ₉₅	Result of Significance Test	0.2	0.7	0.3	0.9	0.3	0.2	0.4	0.3	0.8	1	0.4	2.7	0.4	0.7	0.4	0.4
Arsenic	2	28	12	150	0	250	79.77682	POTENTIALLY SUITABLE FOR USE	72	72		45	49	69	66	82	95		12	16	19	80	21	79
Boron	0.4	28	0.6	3.8	1	3	2.162511	POTENTIALLY SUITABLE FOR USE	2.4	3.8		1	2.8	1.2	1.2	2.2	2.1		0.94	1.3	0.89	1.2	0.88	0.81
Chromium (III)	5	28	31.9	410	1	400	159.8763	POTENTIALLY SUITABLE FOR USE	80	109.85		39	72.85	66	129.9	94	159.9		53.76	41.7	53	219.9	40	89
Chromium (VI)	0.5	19	0.1	0.5	0	25	0.41444	POTENTIALLY SUITABLE FOR USE		0.15			0.15		0.1		0.1		0.24	0.3		0.1		
Copper	5	28	7.3	230	1	200	87.18576	POTENTIALLY SUITABLE FOR USE	25	110		24	230	25	17	30	30		21	15	18	14	9.5	10
Nickel	5	28	23	150	2	110	86.43294	POTENTIALLY SUITABLE FOR USE	81	64		70	77	110	65	91	73		40	29	29	96	24	86
Zinc	10	28	56	410	3	300	229.6131	POTENTIALLY SUITABLE FOR USE	130	260		64	320	120	150	170	180		120	56	110	150	62	220
	Mean																							
pH (su)	7.9				1				7.7	8.1	7.1	8.3	8.1	8.1	8.1	8	8.3	8.3	7.3	7.9	7.9	8.2	7.8	8
Data se	H (su) 7.9 7.9 7.7 8.1 7.1 8.3 8.1 8.1 8.3 8.3 7.3 7.9 7.9 8.2 7.8 8 Risk parameter: Plant life pH >7 Data set: Tip Area (COMBINED) Client: Bovis Barratt and Taylor Wimpey																							

Site: Land at Bankside, Banbury

Job no: C12702







Chemical and data (mg/kg) (blue denotes <= RL)	Mean Abso	STATISTICAL		MENT OF GEO-ENVIRONMENTAL SOIL DATA
(red denotes >= GAC)		= critical value of t		
Beryllium	Potential	Sample	Visual	assessment - Q-q & histogram plots
	Outlier?	•	visual	Q-q plot
1		HTP02 @ 0.20		
2.1		HTP03 @ 0.50		
1.6		HTP04 @ 0.10		3
1.1		HTP05 @ 0.50	_	
0.5		HTP06 @ 0.10 TP1 @ 0.4	_	
0.5		TP1 @ 0.90m		2
0.5		1FT @ 0.30m		
			(i	1
			Sample (yi)	
			<u>e</u>	
			- L	
			- Sai	-3 -2 -1 0 1 2 3
				-1
				-2
		1		-3
		1		Theoretical (zi)
		1		
		1		Beryllium
				2.5
				2
			Number of measurements	
			Ĕ.	1.5
			lre	
			ası	
			ne	
				0.5
			- Q	
				-2 -1 0 1 2 3 4
				Concentration
				Data Normal distribution
			Decia d	Diele novemeter
			Basic d	
			Hur	nan health - residential with plant uptake (2.5%SOM)
				51 = GAC (critical conc.) (mg/kg) $7 = p_0 complex > cr = CAC$
				7 = no. samples $0 = no.$ samples > or = GAC
				0.5 = min. value 2.1 = max. value 1 = laboratory reporting limit (RL)
				2.1 = max. value1 = laboratory reporting limit (RL)571 = mean3 = no. samples at RL
			0.0202	403 = standard deviation RL is limit of detection of the method used
			Ctatiotic	cal tests
				nple t-test One-sided Chebychev Theorem

				$205 = t_0$ $-210.221 = k_0$
				$4.36 = k_{0.05}$ $4.36 = k_{0.05}$
			1.6885	$593 = 95\% \text{ UCL } (\text{US}_{95}) \qquad 2.260837 = 95\% \text{ UCL } (\text{US}_{95})$
				of significance test at 95% confidence level
				thesis (H_0) = level of contamination is the same as, or higher than, the GAC
				we hypotheses (H_1) = level of contamination is lower than the GAC
			Data	a set treated as non-normally distributed
			Therefor	
			Use Ch	ebychev Theorem - Ho rejected, true mean <=GAC
	i			5 = 2.260837 GAC = 51 (US95 = 0.044 x GAC)
		,	Site ref	erence POTENTIALLY SUITABLE FOR USE
			011011011	
			Data set	
				: Tip Area (COMBINED)
			Data set	: Tip Area (COMBINED) Bovis Barratt and Taylor Wimpey
			Data set Client: Site:	: Tip Area (COMBINED) Bovis Barratt and Taylor Wimpey Land at Bankside, Banbury
			Data set	: Tip Area (COMBINED) Bovis Barratt and Taylor Wimpey

Chemical and data

(mg/kg)

(blue denotes <= RL) (red denotes >= GAC)

Boron

0.8 1

1

0.6 1.2

1.6

1.6

2.3

0.86

1.4

2.2 1.7

0.68

2.4

3.8

1

2.8

1.2

1.2

2.2

0.94

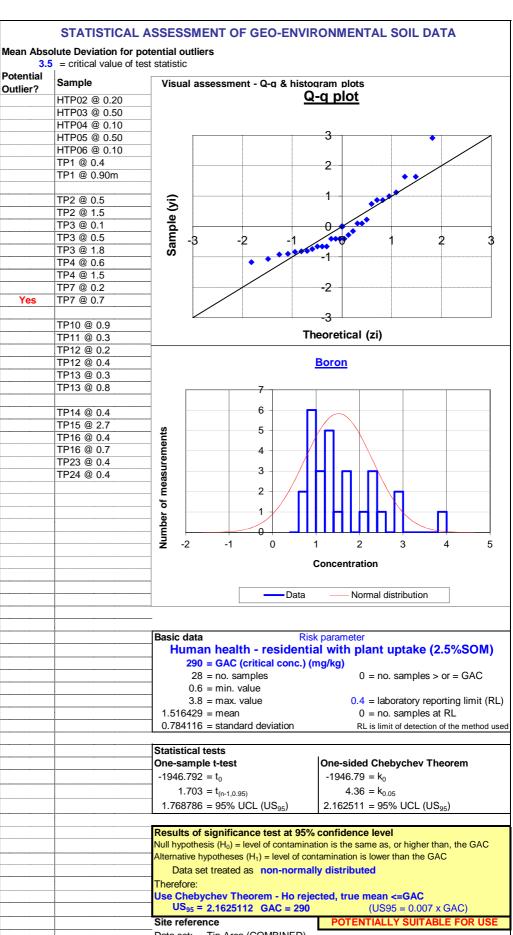
1.3

0.89

1.2

0.88

0.81



Data set treated as non-normall Therefore: Use Chebychev Theorem - Ho reject US ₉₅ = 2.1625112 GAC = 290	∙ sted, true mean <=GAC
Site reference	POTENTIALLY SUITABLE FOR USE
Data set: Tip Area (COMBINED) Client: Bovis Barratt and Taylor V Site: Land at Bankside, Banbur Job no: C12702	
Reference: CL:AIRE & CIEH. May 2008.Gu	idance on comparing soil contamination with a critical concentration.

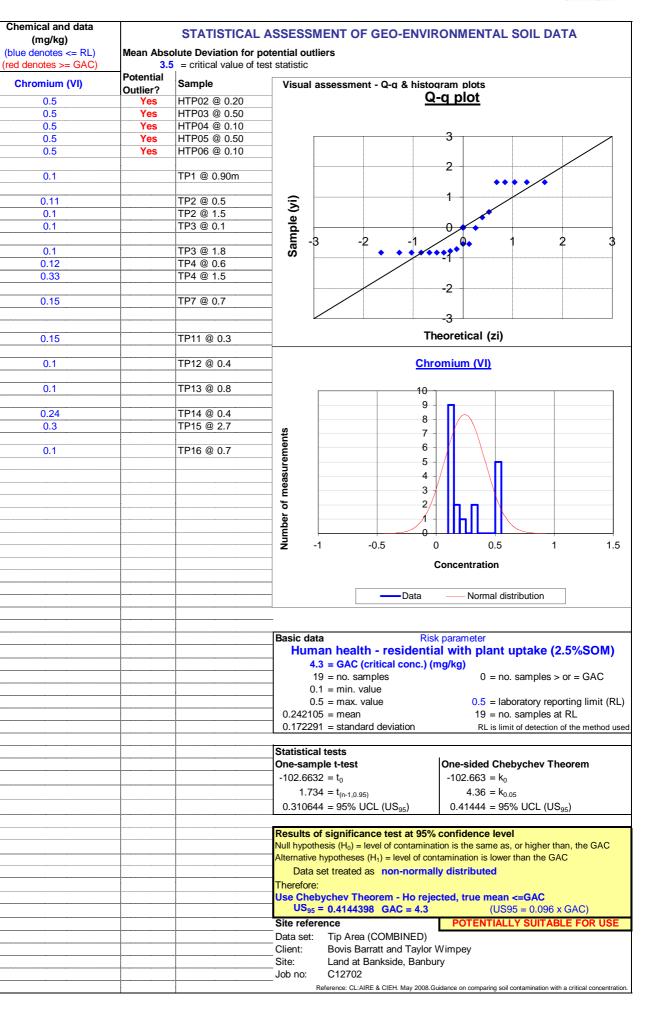




Chemical and data (mg/kg) (blue denotes <= RL) (red denotes >= GAC)		STATISTICAL olute Deviation for p 5 = critical value of t	
	Potential		est statistic Note - MAD Not applicable as 50 % of more of values are the sam
Cadmium	Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.1	n/a	HTP02 @ 0.20	<u>Q-q plot</u>
0.1	n/a	HTP03 @ 0.50	
0.1	n/a	HTP04 @ 0.10	
0.1	n/a	HTP05 @ 0.50	
0.23	n/a	HTP06 @ 0.10	
0.23	n/a	TP1 @ 0.4	
			2
0.5	n/a	TP1 @ 0.90m	
2.5			
0.5	n/a	TP2 @ 0.5	Bamble (Xi)
0.5	n/a	TP2 @ 1.5	
0.5	n/a	TP3 @ 0.1	
0.5	n/a	TP3 @ 0.5	
0.5	n/a	TP3 @ 1.8	S
0.5	n/a	TP4 @ 0.6	
0.5	n/a	TP4 @ 1.5	
0.5	n/a	TP7 @ 0.2	-2
0.52	n/a	TP7 @ 0.7	
0.5	n/a	TP10 @ 0.9	
0.71	n/a	TP11 @ 0.3	Theoretical (zi)
0.5	n/a	TP12 @ 0.2	
0.5	n/a	TP12 @ 0.2	Cadmium
0.5	n/a	TP13 @ 0.3	
0.5	n/a	TP13 @ 0.8	25
0.5	n/a	TP14 @ 0.4	20 1
0.5	n/a	TP15 @ 2.7	
0.5	n/a	TP16 @ 0.4	15 10 10
0.5	n/a	TP16 @ 0.7	
0.5	n/a	TP23 @ 0.4	
0.5	n/a	TP24 @ 0.4	
			5 5
			$ \vec{z}$ -0.5 0 0.5 1 1.5
			Concentration
			Data Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			11 = GAC (critical conc.) (mg/kg)
			28 = no. samples 0 = no. samples > or = GAC
			0.1 = min. value
			0.71 = max. value 0.1 = laboratory reporting limit (RL
			0.445357 = mean 4 = no. samples at RL
			0.159385 = standard deviation RL is limit of detection of the method us
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-350.4089 = t_0 \qquad -350.409 = k_0$
			$1.703 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			0.496653 = 95% UCL (US ₉₅) 0.576684 = 95% UCL (US ₉₅)
			Results of significance test at 95% confidence level
<u> </u>			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses (H_1) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			US ₉₅ = 0.5766844 GAC = 11 (US95 = 0.052 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: Tip Area (COMBINED)
			Client: Bovis Barratt and Taylor Wimpey
			Site: Land at Bankside, Banbury
			Job no: C12702
	1	1	
		1	Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentrati

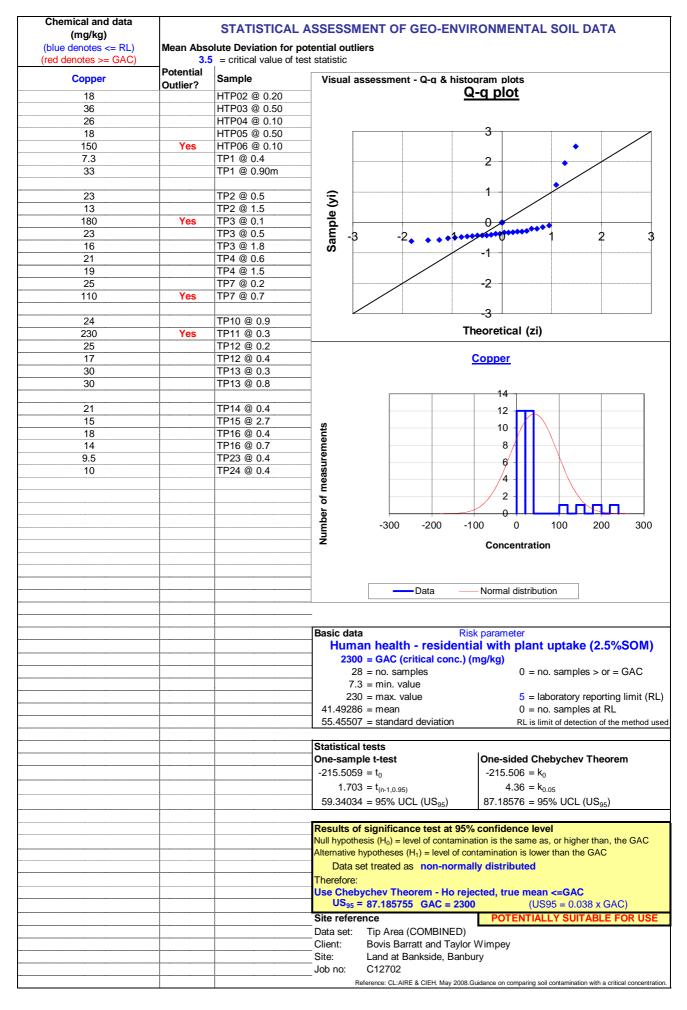


	Chemical and data (mg/kg) (blue denotes <= RL)		olute Deviation for p	potential out		ONMENTAL SOIL DATA
	(red denotes >= GAC)		= critical value of t	est statistic		
	Chromium (III)	Potential Outlier?	Sample	Visual	assessment - Q-q & histoq	ram plots
-	46.5	Outlier	HTP02 @ 0.20		Q-	q plot
	149.5		HTP03 @ 0.50			
	99.5		HTP04 @ 0.10			
	74.5		HTP05 @ 0.50			3
	79.5		HTP06 @ 0.10	_		
-	410	Yes	TP1 @ 0.4	_		2
	159.9		TP1 @ 0.90m			2
	38.89		TP2 @ 0.5	(i		
	31.9		TP2 @ 1.5	Sample (yi)		
	74.9		TP3 @ 0.1	––––––––––––––––––––––––––––––––––––––		
	63		TP3 @ 0.5	<u> </u>	3 -2 -1 🙀	
	129.9		TP3 @ 1.8	Sa		
	62.88		TP4 @ 0.6			
	37.67		TP4 @ 1.5			
	80		TP7 @ 0.2			-2
	109.85		TP7 @ 0.7			
	39		TP10 @ 0.9			-3
	72.85		TP11 @ 0.3		The	eoretical (zi)
	66		TP12 @ 0.2			
	129.9		TP12 @ 0.4		<u>Chro</u>	omium (III)
	94		TP13 @ 0.3			
	159.9		TP13 @ 0.8		8	
					7	
	53.76		TP14 @ 0.4			
	41.7		TP15 @ 2.7	<u>,</u>	6 -	
	53		TP16 @ 0.4	ent	5 -	
	219.9		TP16 @ 0.7	Number of measurements		
	40		TP23 @ 0.4	nre	4 -	
	89		TP24 @ 0.4	as	3 -	
				e		
					2/-	
				er	1-	
				- de	0	
				2	-400 -200 0	200 400 600
					C	Concentration
					Data	Normal distribution
				unununununu		
				Decia de	te Diele	
				Basic da		parameter
						I with plant uptake (2.5%SOM)
					30 = GAC (critical conc.) (m	
					28 = no. samples	0 = no. samples > or = GAC
					.9 = min. value	
					10 = max. value	5 = laboratory reporting limit (RL)
					86 = mean	0 = no. samples at RL
				76.682	42 = standard deviation	RL is limit of detection of the method used
				Statiati-	al tooto	
				Statistic		One sided Chebyshey Theorem
					•	One-sided Chebychev Theorem
				-36.801	-	$-36.8011 = k_0$
				1.7	$03 = t_{(n-1,0.95)}$	$4.36 = k_{0.05}$
1				121.37	21 = 95% UCL (US ₉₅)	159.8763 = 95% UCL (US ₉₅)
		1				
				Results	of significance test at 95%	confidence level
-				Null hypot	hesis (H ₀) = level of contaminati	on is the same as, or higher than, the GAC
				Alternative	hypotheses (H1) = level of cont	amination is lower than the GAC
				Data	set treated as non-normally	y distributed
				Therefore):	
					bychev Theorem - Ho rejec	cted, true mean <=GAC
		1			= 159.8763 GAC = 630	(US95 = 0.254 x GAC)
				Site refe	rence	POTENTIALLY SUITABLE FOR USE
				Data set:		
			1	Client:	Bovis Barratt and Taylor V	Vimpev
				Site:	Land at Bankside, Banbur	
				J		1
				Joh no:	C12702	
				Job no:	C12702 Reference: CL:AIRE & CIEH. May 2008.Gu	- idance on comparing soil contamination with a critical concentration.



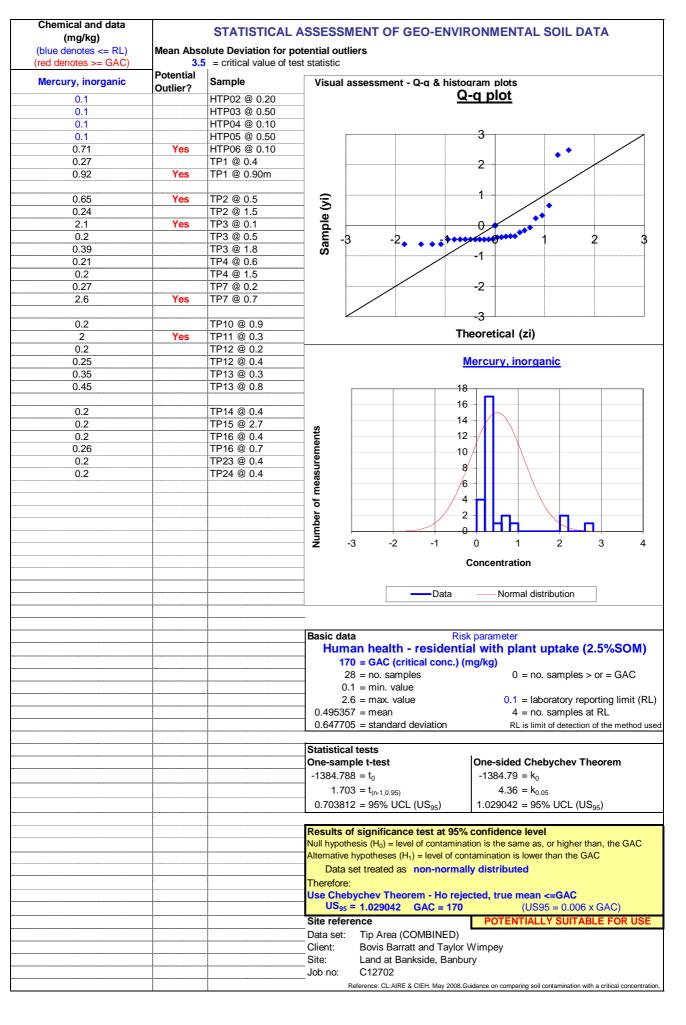








Chemical and data (mg/kg) (blue denotes <= RL) (red denotes >= GAC)		STATISTICAL blute Deviation for p = critical value of te	
	Potential		
Lead	Outlier?	Sample	Visual assessment - Q-q & histogram plots
39		HTP02 @ 0.20	<u>Q-q plot</u>
230	Yes	HTP03 @ 0.50	
52		HTP04 @ 0.10	3
<u> </u>	Vac	HTP05 @ 0.50 HTP06 @ 0.10	
400	Yes	TP1 @ 0.4	
75		TP1 @ 0.90m	2
62		TP2 @ 0.5	
17		TP2 @ 1.5	
410	Yes	TP3 @ 0.1	Sample (i)
24		TP3 @ 0.5	
33		TP3 @ 1.8	/Ő
51		TP4 @ 0.6	
<u> </u>		TP4 @ 1.5 TP7 @ 0.2	
360	Yes	TP7 @ 0.2 TP7 @ 0.7	
500	105		
29		TP10 @ 0.9	
1500	Yes	TP11 @ 0.3	Theoretical (zi)
40		TP12 @ 0.2	
40		TP12 @ 0.4	Lead
74		TP13 @ 0.3	
81		TP13 @ 0.8	
			25
23		TP14 @ 0.4	20
22		TP15 @ 2.7	
<u>21</u> 22		TP16 @ 0.4 TP16 @ 0.7	15/10 10
19		TP23 @ 0.4	
25		TP24 @ 0.4	
20		11 24 8 0.4	
			-1500 -1000 -500 0 500 1000 1500 2000
			<u> </u>
			Z Concentration
			Data Normal distribution
			Desis data Disk sovereter
			Basic data Risk parameter Human health - residential with plant uptake (2.5%SOM)
			450 = GAC (critical conc.) (mg/kg) 28 = no. samples 2 = no. samples > or = GAC
			28 = no. samples 2 = no. samples > or = GAC 17 = min. value
			1500 = max. value 5 = laboratory reporting limit (RL)
			$138.6786 = mean \qquad \qquad 0 = no. samples at RL$
			294.188 = standard deviation RL is limit of detection of the method us
		1	
			Statistical tests
		1	One-sample t-test One-sided Chebychev Theorem
		1	$-5.599678 = t_0$ $-5.59968 = k_0$
		1	$1.703 = t_{(n-1,0.95)} 4.36 = k_{0.05}$
			$233.3591 = 95\% \text{ UCL } (\text{US}_{95}) \qquad 381.0785 = 95\% \text{ UCL } (\text{US}_{95})$
			Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
		1	
			Alternative hypotheses (H ₁) = level of contamination is lower than the GAC
			Alternative hypotheses (H ₁) = level of contamination is lower than the GAC Data set treated as non-normally distributed
			Data set treated as non-normally distributed
			Data set treated as non-normally distributed Therefore:
			Data set treated as non-normally distributed
			Data set treated as non-normally distributed Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			Data set treated as non-normally distributed Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			Data set treated as non-normally distributed Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			Data set treated as non-normally distributed Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			Data set treated as non-normally distributed Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC





Chemical and data

(mg/kg)

(blue denotes <= RL) (red denotes >= GAC)

Nickel

25

64 67

45

53 150

80

23

28 52

80

59

31

33

81

64

70

77

110

65 91 Mean Absolute Deviation for potential outliers 3.5 = critical value of test statistic

Sample

HTP02 @ 0.20

HTP03 @ 0.50

HTP04 @ 0.10 HTP05 @ 0.50

HTP06 @ 0.10

TP1 @ 0.90m

Sample (yi)

-3

TP1 @ 0.4

TP2 @ 0.5

TP2 @ 1.5

TP3 @ 0.1

TP3 @ 0.5

TP3 @ 1.8

TP4 @ 0.6

TP4 @ 1.5 TP7 @ 0.2

TP7 @ 0.7

TP10 @ 0.9

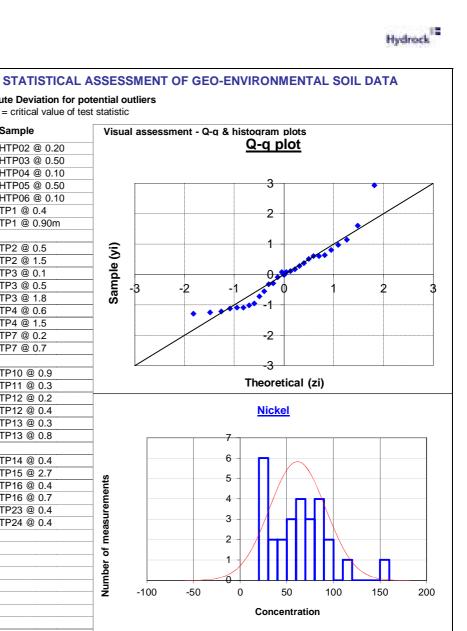
TP11 @ 0.3 TP12 @ 0.2

TP12 @ 0.4

TP13 @ 0.3

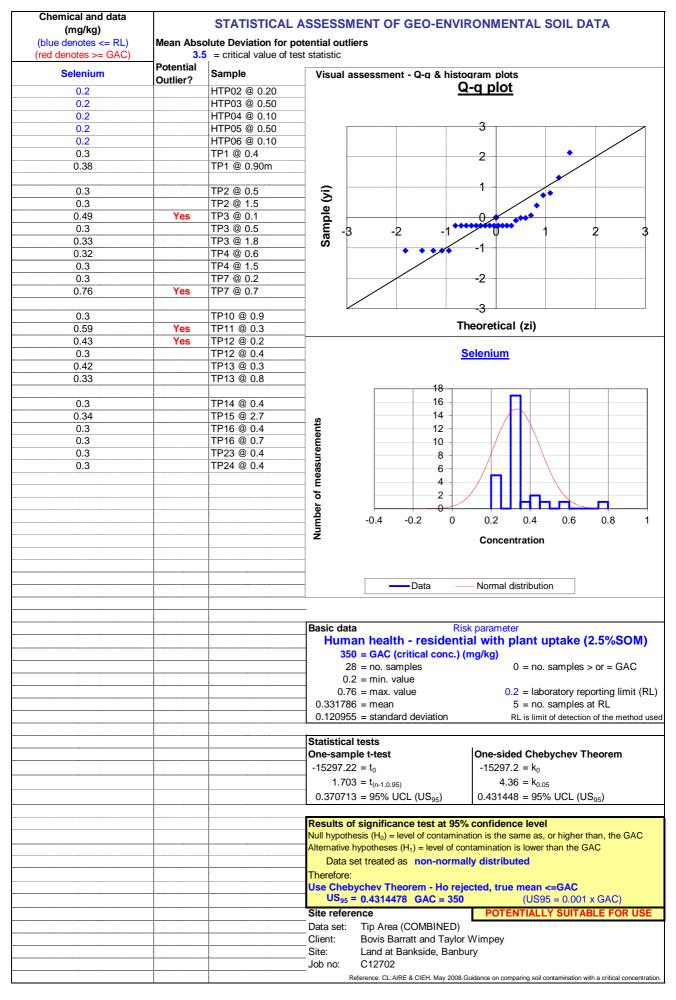
Potential

Outlier?



70	IP13 @ 0.3	
73	TP13 @ 0.8	7
40	TP14 @ 0.4	
29	TP15 @ 2.7	
29	TP16 @ 0.4	steer st
96	TP16 @ 0.7	
24	TP23 @ 0.4	
86	TP24 @ 0.4	
		¹ ¹ ¹ ¹ ¹ ¹ ¹ ¹ ¹
		ž -100 -50 0 50 100 150 200
		Concentration
		Data Normal distribution
		Basic data Risk parameter
		Human health - residential with plant uptake (2.5%SOM)
		130 = GAC (critical conc.) (mg/kg)
		28 = no. samples 1 = no. samples > or = GAC
	1	23 = min. value
		150 = max. value 5 = laboratory reporting limit (R
		61.60714 = mean $0 = no. samples at RL$
		30.12977 = standard deviation RL is limit of detection of the method u
		Statistical tests
		One-sample t-test One-sided Chebychev Theorem
		One-sample t-test One-sided Chebychev Theorem -12.01141 = t ₀ -12.0114 = k ₀
		One-sample t-test One-sided Chebychev Theorem -12.01141 = t ₀ -12.0114 = k ₀ 1.703 = t _(n-1,0.95) 4.36 = k _{0.05}
		One-sample t-test One-sided Chebychev Theorem -12.01141 = t ₀ -12.0114 = k ₀
		$\begin{tabular}{ c c c c c c c c c c c c c c c c c c c$
		$\begin{tabular}{ c c c c c c c c c c c c c c c c c c c$
		$\begin{tabular}{ c c c c c c c c c c c c c c c c c c c$
		$\begin{tabular}{ c c c c c c c c c c c c c c c c c c c$
		$\label{eq:constraint} \begin{array}{ c c c } \hline \textbf{One-sided Chebychev Theorem} \\ -12.01141 = t_0 & -12.0114 = k_0 \\ 1.703 = t_{(n-1,0.95)} & 4.36 = k_{0.05} \\ \hline 71.30401 = 95\% \ \text{UCL} \ (\text{US}_{95}) & 86.43294 = 95\% \ \text{UCL} \ (\text{US}_{95}) \\ \hline \textbf{Results of significance test at 95\% confidence level} \\ \hline \text{Null hypothesis} \ (H_0) = \text{level of contamination is the same as, or higher than, the GAC} \\ \hline \text{Alternative hypotheses} \ (H_1) = \text{level of contamination is lower than the GAC} \\ \hline \text{Data set treated as non-normally distributed} \\ \hline \end{array}$
		$\label{eq:constraint} \begin{array}{ c c c } \hline \textbf{One-sample t-test} & \textbf{One-sided Chebychev Theorem} \\ -12.01141 = t_0 & -12.0114 = k_0 \\ 1.703 = t_{(n-1,0.95)} & 4.36 = k_{0.05} \\ \hline 71.30401 = 95\% \ \text{UCL} \ (\text{US}_{95}) & 86.43294 = 95\% \ \text{UCL} \ (\text{US}_{95}) \\ \hline \hline \textbf{Results of significance test at 95\% confidence level} \\ \hline \text{Null hypothesis} \ (H_0) = \text{level of contamination is the same as, or higher than, the GAC} \\ \hline \text{Alternative hypotheses} \ (H_1) = \text{level of contamination is lower than the GAC} \\ \hline \text{Data set treated as non-normally distributed} \\ \hline \hline \textbf{Therefore:} \end{array}$
		One-sample t-testOne-sided Chebychev Theorem $-12.01141 = t_0$ $-12.0114 = k_0$ $1.703 = t_{(n-1,0.95)}$ $4.36 = k_{0.05}$ $71.30401 = 95\%$ UCL (US ₉₅) $86.43294 = 95\%$ UCL (US ₉₅)Results of significance test at 95% confidence levelNull hypothesis (H ₀) = level of contamination is the same as, or higher than, the GACAlternative hypotheses (H ₁) = level of contamination is lower than the GACData set treated as non-normally distributedTherefore:Use Chebychev Theorem - Ho rejected, true mean <=GAC
		$\label{eq:constraints} \begin{array}{llllllllllllllllllllllllllllllllllll$
		One-sample t-testOne-sided Chebychev Theorem $-12.01141 = t_0$ $-12.0114 = k_0$ $1.703 = t_{(n-1,0.95)}$ $4.36 = k_{0.05}$ $71.30401 = 95\%$ UCL (US ₉₅) $86.43294 = 95\%$ UCL (US ₉₅)Results of significance test at 95% confidence levelNull hypothesis (H ₀) = level of contamination is the same as, or higher than, the GACAlternative hypotheses (H ₁) = level of contamination is lower than the GACData set treated as non-normally distributedTherefore:Use Chebychev Theorem - Ho rejected, true mean <=GAC
		$\label{eq:constraints} \begin{array}{llllllllllllllllllllllllllllllllllll$
		One-sample t-testOne-sided Chebychev Theorem $-12.01141 = t_0$ $-12.0114 = k_0$ $1.703 = t_{(n-1,0.95)}$ $4.36 = k_{0.05}$ $71.30401 = 95\%$ UCL (US ₉₅) $86.43294 = 95\%$ UCL (US ₉₅)Results of significance test at 95% confidence levelNull hypothesis (H ₀) = level of contamination is the same as, or higher than, the GACAlternative hypotheses (H ₁) = level of contamination is lower than the GACData set treated as non-normally distributedTherefore:Use Chebychev Theorem - Ho rejected, true mean <=GAC
		One-sample t-testOne-sided Chebychev Theorem $-12.01141 = t_0$ $-12.0114 = k_0$ $1.703 = t_{(n-1,0.95)}$ $4.36 = k_{0.05}$ $71.30401 = 95\%$ UCL (US ₉₅) $86.43294 = 95\%$ UCL (US ₉₅)Results of significance test at 95% confidence levelNull hypothesis (H_0) = level of contamination is the same as, or higher than, the GACAlternative hypotheses (H_1) = level of contamination is lower than the GACData set treated as non-normally distributedTherefore:Use Chebychev Theorem - Ho rejected, true mean <=GAC
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		One-sample t-testOne-sided Chebychev Theorem $-12.01141 = t_0$ $-12.0114 = k_0$ $1.703 = t_{(n-1,0.95)}$ $4.36 = k_{0.05}$ $71.30401 = 95\%$ UCL (US ₉₅) $86.43294 = 95\%$ UCL (US ₉₅)Results of significance test at 95% confidence levelNull hypothesis (H_0) = level of contamination is the same as, or higher than, the GACAlternative hypotheses (H_1) = level of contamination is lower than the GACData set treated as non-normally distributedTherefore:Use Chebychev Theorem - Ho rejected, true mean <=GAC



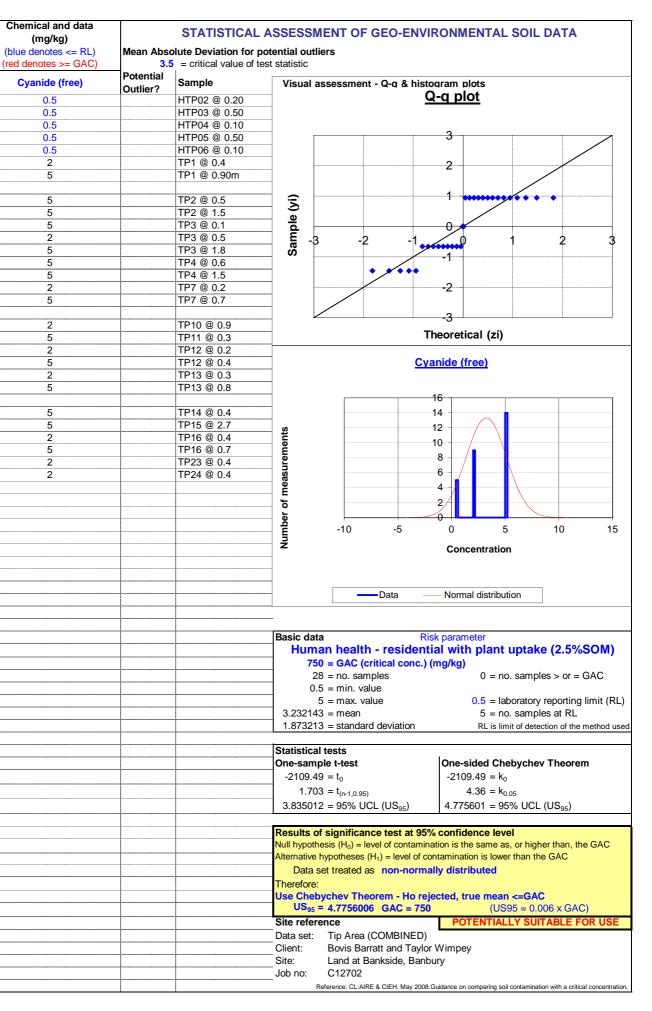




Chemical and data (mg/kg)		STATISTICAL	ASSES	SMENT OF GEO-ENVIR	ONMENTAL SOIL DATA
(blue denotes <= RL) (red denotes >= GAC)	3.5	blute Deviation for p = critical value of te	otential o	utliers	
Vanadium	Potential Outlier?	Sample	Visua	al assessment - Q-q & histor	
67		HTP02 @ 0.20	_	<u>Q</u> .	<u>-q plot</u>
230		HTP03 @ 0.50			
150		HTP04 @ 0.10			
95		HTP05 @ 0.50			3
120		HTP06 @ 0.10			
0.5		TP1 @ 0.4	_		2
0.5		TP1 @ 0.90m			-
			Ē		1
			Sample (yi)		0
			E	-3 -2 -1	🗡 0 1 2 3
			ů		_1
				•	
					-2
					-3
				Th	eoretical (zi)
				111	
					modium
				Va	anadium
				4.	5
					4
				3.	
			t		3 - / / / / / / / / / / / / / / / / / /
				2.	
			e u		
			Number of measurements		
			ea	1.	5
			E		
			ō	0.	
			pe		
			<u> </u>	-400 -200	0 200 400 600
· · ·			Ī		Concentration
· · ·			-		
				_	
				Data	Normal distribution
			Basic		parameter
			Hu		al with plant uptake (2.5%SOM)
				74 = GAC (critical conc.) (n	
				7 = no. samples	4 = no. samples > or = GAC
				0.5 = min. value	_
				230 = max. value	5 = laboratory reporting limit (RL)
				429 = mean (mean>GAC)	2 = no. samples at RL
		1	82.17	758 = standard deviation	RL is limit of detection of the method used
			Ctarl.	ical tests	
		+		ical tests	One-sided Chebyshov Theorem
				ample t-test	One-sided Chebychev Theorem 0.666908 = k ₀
				$3908 = t_0$	0
· · · · · · · · · · · · · · · · · · ·				$.943 = t_{(n-1,0.95)}$	$4.36 = k_{0.05}$
			155.0	643 = 95% UCL (US ₉₅)	230.1368 = 95% UCL (US ₉₅)
				s of significance test at 95%	
					ion is the same as, or higher than, the GAC
				ive hypotheses $(H_1) =$ level of cont	
				ta set treated as non-normall	y distributed
			Therefo		
				hebychev Theorem - Ho acce	
				95 = 230.13678 GAC = 74	(US95 = 3.11 x GAC)
				ference	FURTHER ASSESSMENT REQUIRED
			Data se	1 (,	
			Client:	Bovis Barratt and Taylor V	
			Site:	Land at Bankside, Banbu	ry
			Job no	C12702	
		Transmoot in the second s		Reference: CL:AIRE & CIEH. May 2008.Gu	idance on comparing soil contamination with a critical concentration.



Chemical and data (mg/kg) (blue denotes <= RL) (red denotes >= GAC)		STATISTICAL blute Deviation for p = critical value of t	potential ou	MENT OF GEO-ENVIRONMENTAL SOIL DATA
	Potential			
Zinc	Outlier?	Sample	Visual	assessment - Q-q & histogram plots
79		HTP02 @ 0.20		<u>Q-q plot</u>
170		HTP03 @ 0.50		
190		HTP04 @ 0.10		0
110		HTP05 @ 0.50		3
410	Yes	HTP06 @ 0.10		
260		TP1 @ 0.4		2
180		TP1 @ 0.90m		
			_	
130		TP2 @ 0.5	Sample (yi)	
64		TP2 @ 1.5	<u> </u>	
300		TP3 @ 0.1	d	
59		TP3 @ 0.5	- a -	β -2 -1 -1 -2 -3
160		TP3 @ 1.8	S	-1
<u>98</u> 64		TP4 @ 0.6		
130		TP4 @ 1.5 TP7 @ 0.2		
260		TP7 @ 0.2		
260		IP7 @ 0.7		
61		TP10 @ 0.9		-3
<u> </u>		TP10 @ 0.9 TP11 @ 0.3		Theoretical (zi)
		TP11 @ 0.3		
<u> </u>		TP12 @ 0.2 TP12 @ 0.4		Zinc
150		TP12 @ 0.4 TP13 @ 0.3		Zinc
170		TP13 @ 0.3		
100		1 F 13 @ U.O		6
120		TP14 @ 0.4		
56		TP15 @ 2.7		5 - 1
110		TP16 @ 0.4	t	4
150		TP16 @ 0.7	measurements	
62		TP23 @ 0.4	Len	3 3
220		TP24 @ 0.4		
220			lea	
· · · · · · · · · · · · · · · · · · ·			E	
			ō	
			Number of	
			Ę	-400 -200 0 200 400 600
			ž	Concentration
· · · ·			-	
		-		
				Data Normal distribution
			Basic da	ta Risk parameter
			Hun	nan health - residential with plant uptake (2.5%SOM)
			37	00 = GAC (critical conc.) (mg/kg)
				28 = no. samples $0 = no. samples > or = GAC$
				56 = min. value
			4	10 = max. value 10 = laboratory reporting limit (RL)
			156.64	29 = mean 0 = no. samples at RL
			88.560	18 = standard deviation RL is limit of detection of the method use
			Statistic	
			One-sar	nple t-test One-sided Chebychev Theorem
			-211.71	
		1		$03 = t_{(n-1,0.95)}$ $4.36 = k_{0.05}$
		1		$48 = 95\% \text{ UCL } (\text{US}_{95}) $ $229.6131 = 95\% \text{ UCL } (\text{US}_{95})$
			100.14	223.0101 - 30 % OOL (0095)
			Perult	of significance test at 05% confidence lavel
				of significance test at 95% confidence level thesis (H_0) = level of contamination is the same as, or higher than, the GAC
				the size $(H_0) = 1000$ for contamination is the same as, or higher than, the GAC by hypotheses $(H_1) = 1000$ for contamination is lower than the GAC
				set treated as non-normally distributed
			Therefor	
				bychev Theorem - Ho rejected, true mean <=GAC
				EXAMPLE 229.61313 GAC = 3700 (US95 = 0.062 x GAC)
			Site refe	
			Data set	
		ļ	Client:	Bovis Barratt and Taylor Wimpey
	1	1	Site:	Land at Bankside, Banbury
				-
			Job no:	C12702







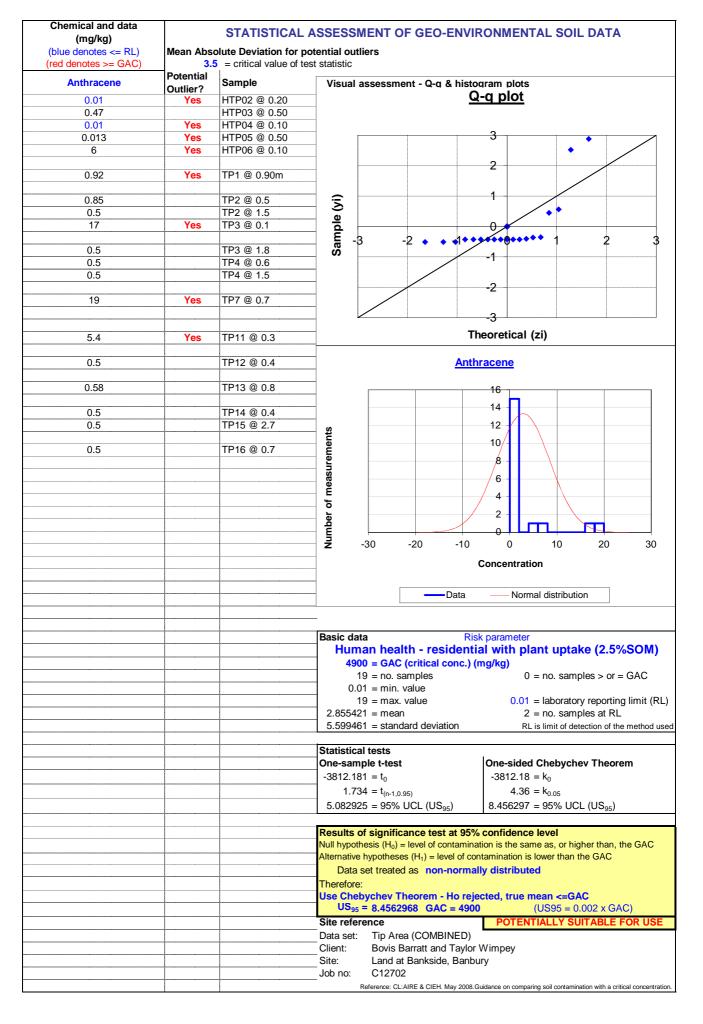
(mg/kg) ue denotes <= RL) denotes >= GAC)		STATISTICAL olute Deviation for p 5 = critical value of t	
 ,	Potential		ssi statistic Note - MAD hot applicable as 50% of more of values are the same
 Phenol (total)	Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.3	n/a	HTP02 @ 0.20	<u>Q-q plot</u>
 0.3	n/a	HTP03 @ 0.50	
 0.3	n/a	HTP04 @ 0.10	
 0.3	n/a	HTP05 @ 0.50	3
 0.3	n/a	HTP06 @ 0.10	
 0.5	n/a	TP1 @ 0.4	
 4.3	n/a	TP1 @ 0.90m	
 0.5	n/a	TP2 @ 0.5	
 0.5	n/a	TP2 @ 1.5	
 3.7	n/a	TP3 @ 0.1	
 0.5	n/a	TP3 @ 0.5	
 0.5	n/a	TP3 @ 1.8	
 0.5	n/a	TP4 @ 0.6	
 0.5	n/a	TP4 @ 1.5	
 0.5	n/a	TP7 @ 0.2	-2
 4.8	n/a	TP7 @ 0.7	
 0.5	n/a	TP10 @ 0.9	
 2.1	n/a	TP11 @ 0.3	Theoretical (zi)
 0.5	n/a	TP12 @ 0.2	
 0.5	n/a	TP12 @ 0.4	Phenol (total)
 0.5	n/a	TP13 @ 0.3	-
 0.58	n/a	TP13 @ 0.8	
 0.5	n/a	TP14 @ 0.4	
 0.76	n/a	TP15 @ 2.7	
 		TP16 @ 0.4	
 0.5	n/a	TP16 @ 0.4	stremen b and b a a a a a a a a a a a a a a a a a a
 0.5	n/a		
 0.5	n/a	TP23 @ 0.4	
 0.5	n/a	TP24 @ 0.4	
 			- To
 			
			5
			Z Concentration
 			Data Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
 			290 = GAC (critical conc.) (mg/kg)
			28 = no. samples $0 = no. samples > or = GAC$
 			0.3 = min. value
 			4.8 = max. value $0.3 = laboratory reporting limit (RL)$
 			0.937143 = mean $5 = no. samples at RL$
 			1.227062 = standard deviation RL is limit of detection of the method use
 			RL IS IMIT OF DETECTION OF THE METHOD USE
 			Statistical tests
 			One-sample t-test One-sided Chebychev Theorem
			$-1246.536 = t_0$ $-1246.54 = k_0$
			$1.703 = t_{(n-1,0.95)}$ $4.36 = k_{0.05}$
 · · ·			1.332056 = 95% UCL (US ₉₅) $1.948196 = 95%$ UCL (US ₉₅)
 		1	Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
 			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H_1) = level of contamination is lower than the GAC
 			Null hypothesis (H ₀) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H ₁) = level of contamination is lower than the GAC Data set treated as non-normally distributed
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H_1) = level of contamination is lower than the GAC Data set treated as non-normally distributed Therefore:
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H_1) = level of contamination is lower than the GAC Data set treated as non-normally distributed Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			Null hypothesis (H ₀) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H ₁) = level of contamination is lower than the GAC Data set treated as non-normally distributed Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H_1) = level of contamination is lower than the GAC Data set treated as non-normally distributed Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			Null hypothesis (H ₀) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H ₁) = level of contamination is lower than the GAC Data set treated as non-normally distributed Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			Null hypothesis (H ₀) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H ₁) = level of contamination is lower than the GAC Data set treated as non-normally distributed Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			Null hypothesis (H ₀) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H ₁) = level of contamination is lower than the GAC Data set treated as non-normally distributed Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC



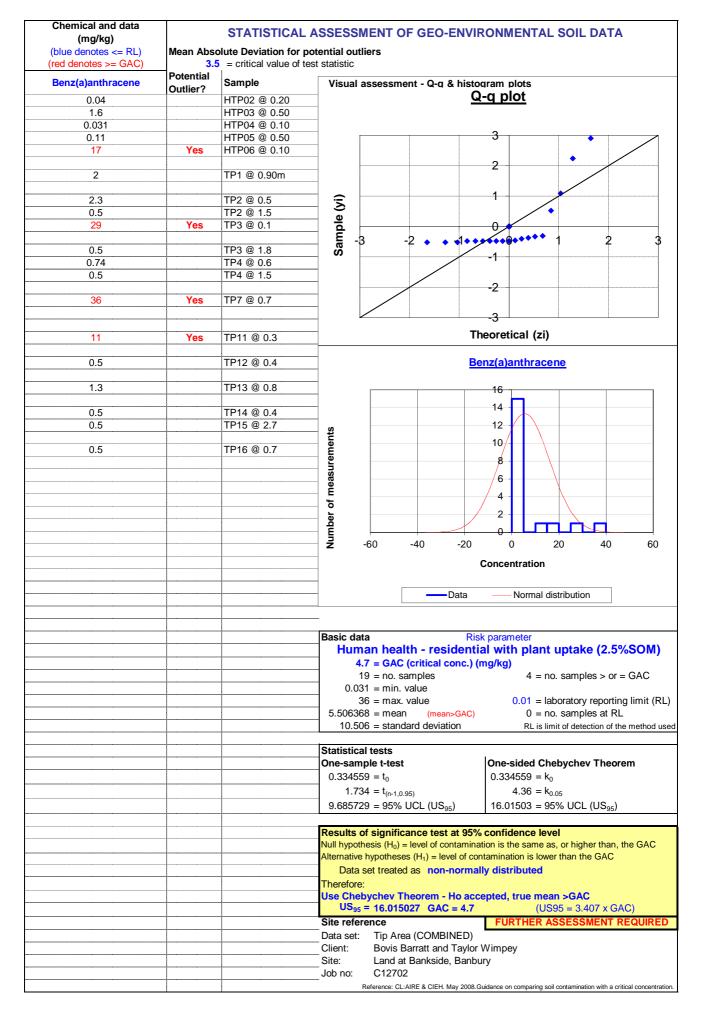
	Chemical and data (mg/kg) (blue denotes <= RL)		olute Deviation for p	
	(red denotes >= GAC)		= critical value of to	est statistic Note - MAD not applicable as 50% or more of values are the same.
	Acenaphthene	Potential	Sample	Visual assessment - Q-q & histogram plots
		Outlier?	•	Q-q plot
	0.01	n/a	HTP02 @ 0.20	
	0.088	n/a	HTP03 @ 0.50	
	0.01	n/a	HTP04 @ 0.10	
	0.01	n/a	HTP05 @ 0.50	3
	0.73	n/a	HTP06 @ 0.10	
	0.5	n/a	TP1 @ 0.90m	2
	0 F	n/a	TP2 @ 0.5	
	0.5	n/a		
		n/a	TP2 @ 1.5	<u>e</u>
	0.5	n/a	TP3 @ 0.1	
	0.5	n/a	TP3 @ 1.8	
	0.5	n/a	TP4 @ 0.6	-1-1
	0.5	n/a	TP4 @ 0.0	
	0.5	11/a	1P4 @ 1.5	
	11	n/a	TP7 @ 0.7	
	2.7	n/a	TP11 @ 0.3	Theoretical (zi)
	0.5	n/a	TP12 @ 0.4	Acenaphthene
	0.5	n/a	TP13 @ 0.8	
	0.5	n/a	TP14 @ 0.4	
	0.5	n/a n/a	TP14 @ 0.4 TP15 @ 2.7	14
	0.5	n/a	1115 @ 2.7	
	0.5	n/a	TP16 @ 0.7	
				-10 -5 0 5 10 15 Concentration
				Data — Normal distribution
				Basic data Risk parameter
				Human health - residential with plant uptake (2.5%SOM)
				480 = GAC (critical conc.) (mg/kg)
				19 = no. samples $0 = no. samples > or = GAC$
				0.01 = min. value
				11 = max. value $0.01 = laboratory reporting limit (RL)$
				1.081474 = mean $3 = no. samples at RL$
				2.467667 = standard deviation RL is limit of detection of the method used
				Statistical tests
				One-sample t-test One-sided Chebychev Theorem
				$-845.9641 = t_0 \qquad -845.964 = k_0$
				$1.734 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
				2.063128 = 95% UCL (US ₉₅) $3.549764 = 95%$ UCL (US ₉₅)
				Results of significance test at 95% confidence level
				Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
				Alternative hypotheses (H ₁) = level of contamination is lower than the GAC Data set treated as non-normally distributed
				Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
				$US_{95} = 3.5497637 \text{ GAC} = 480 $ (US95 = 0.007 x GAC)
				Site reference POTENTIALLY SUITABLE FOR USE
				Data set: Tip Area (COMBINED)
				Client: Bovis Barratt and Taylor Wimpey
				Site: Land at Bankside, Banbury
				Job no: C12702
1		-		Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.



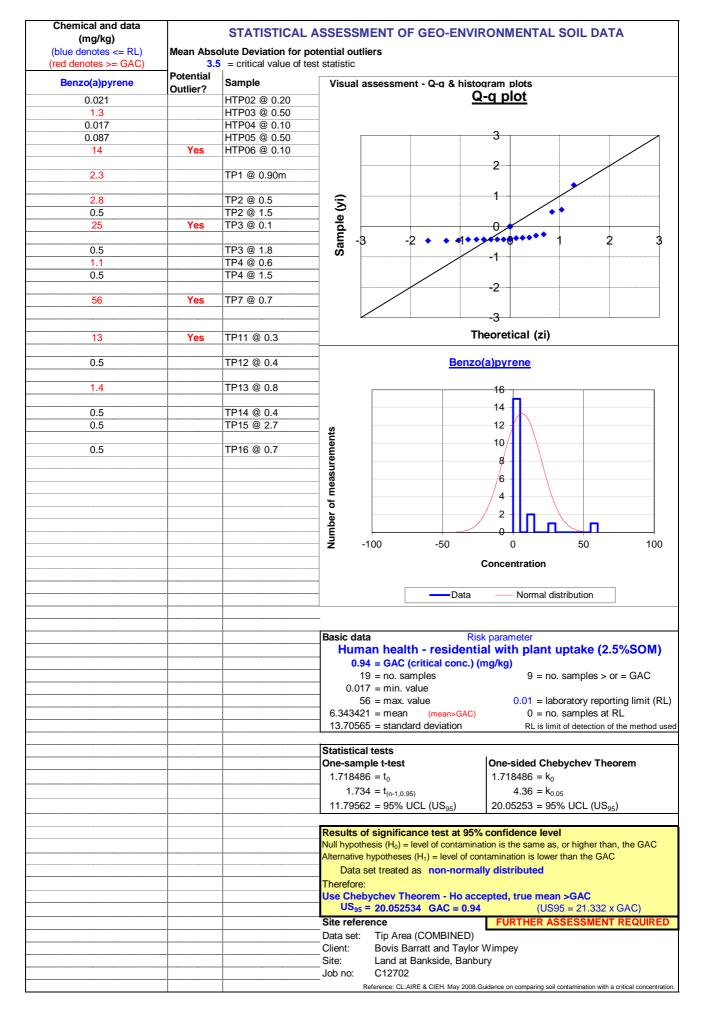
Chemical and data (mg/kg) (blue denotes <= RL) (red denotes >= GAC)		STATISTICAL olute Deviation for p = critical value of to	otential ou	Itliers	
				Note - MAD not applica	able as 50% or more of values are the same.
Acenaphthylene	Outlier?	Sample	Visua	l assessment - Q-q & histoq	
0.01	n/a	HTP02 @ 0.20		<u>Q-</u>	<u>-q plot</u>
0.2	n/a	HTP03 @ 0.50			
0.01	n/a	HTP04 @ 0.10			2
0.01	n/a	HTP05 @ 0.50	_		3
2.8	n/a	HTP06 @ 0.10			2
0.5	n/a	TP1 @ 0.90m			1
0.5	n/a	TP2 @ 0.5	<u> </u>		
0.5	n/a	TP2 @ 1.5	<u> </u>		
5.4	n/a	TP3 @ 0.1	Sample (yi)	-3 -2 -1-	
0.5	n/a	TP3 @ 1.8	Sa		1
0.5	n/a	TP4 @ 0.6			-1
0.5	n/a	TP4 @ 1.5			-2
1.5	n/a	TP7 @ 0.7			-3
0.51	n/a	TP11 @ 0.3		The	eoretical (zi)
0.5	n/a	TP12 @ 0.4		Acen	aphthylene
0.5	n/a	TP13 @ 0.8		14	
				12	
0.5	n/a n/a	TP14 @ 0.4 TP15 @ 2.7	s	12	
0.5	n/a	TP16 @ 0.7	ement	8	
			Number of measurements	6	
				2	
			Nun	-6 -4 -2	0 2 4 6 8
				с	Concentration
				Data	Normal distribution
			Basic d		parameter
					I with plant uptake (2.5%SOM)
			4	<pre>100 = GAC (critical conc.) (m 19 = no. samples</pre>	1g/kg) 0 = no. samples > or = GAC
			0	.01 = min. value	0 = 10. samples > $01 = GAC$
				5.4 = max. value	0.01 = laboratory reporting limit (RL)
				947 = mean	3 = no. samples at RL
			1.267	771 = standard deviation	RL is limit of detection of the method used
				cal tests	One sided Chalumber Th
				mple t-test 177 = t ₀	One-sided Chebychev Theorem -1372.48 = k ₀
				$734 = t_{(n-1,0.95)}$	$4.36 = k_{0.05}$
				251 = 95% UCL (US ₉₅)	2.106977 = 95% UCL (US ₉₅)
			Results	of significance test at 95%	confidence level
			Null hypo	othesis $(H_0) =$ level of contaminati	ion is the same as, or higher than, the GAC tamination is lower than the GAC
			Data	a set treated as non-normall	
				ebychev Theorem - Ho rejec	cted, true mean <=GAC
			USg	₅ = 2.1069772 GAC = 400	(US95 = 0.005 x GAC)
			Data set		POTENTIALLY SUITABLE FOR USE
			Client:	Bovis Barratt and Taylor V	Vimpey
			Site:	Land at Bankside, Banbur	
			Job no:	C12702	
				Reference: CL:AIRE & CIEH. May 2008.Gu	idance on comparing soil contamination with a critical concentration.
•		•			



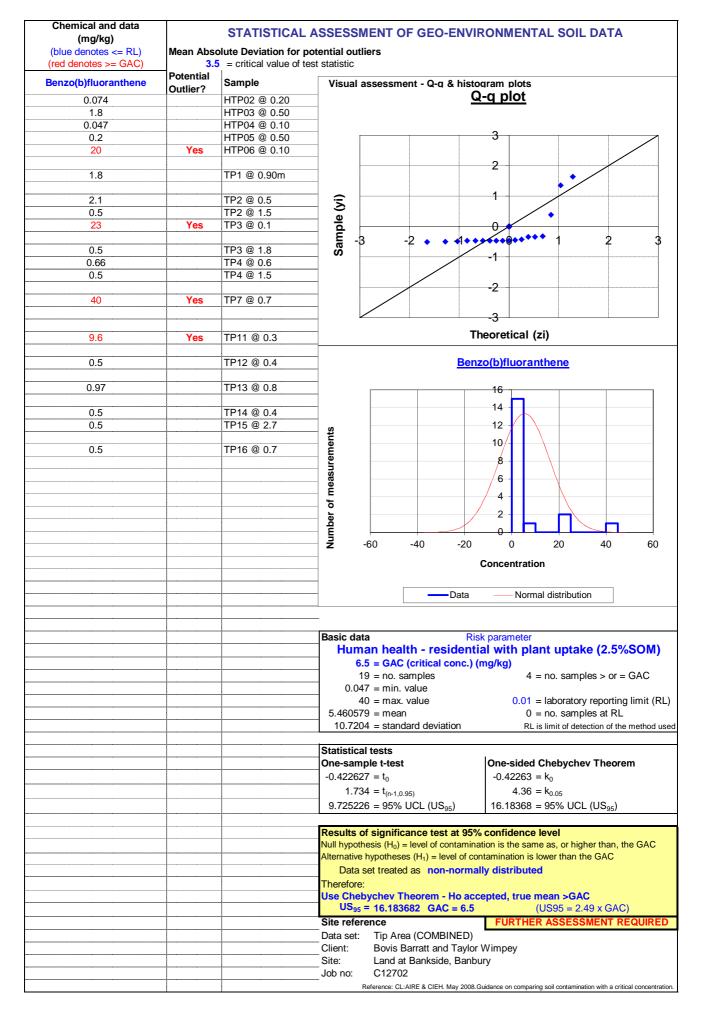




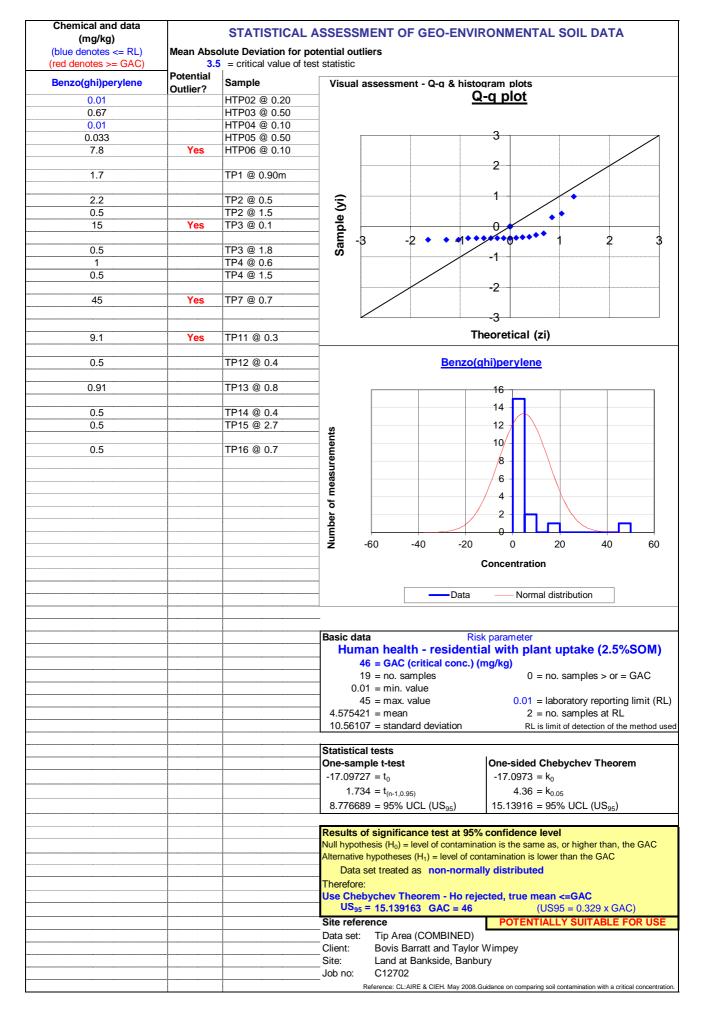




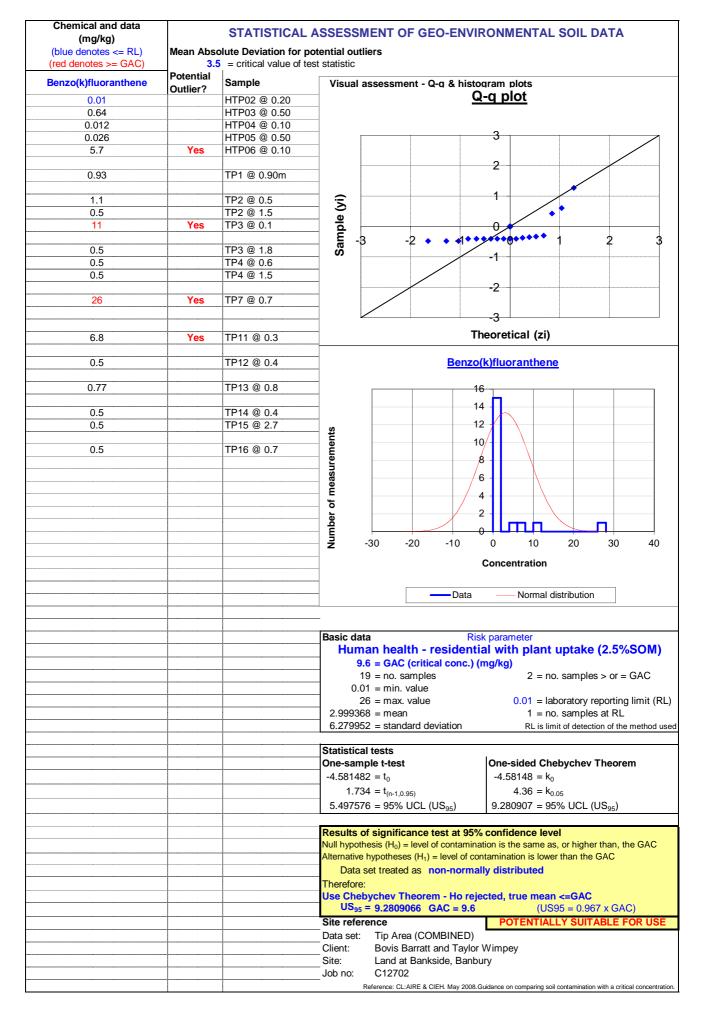




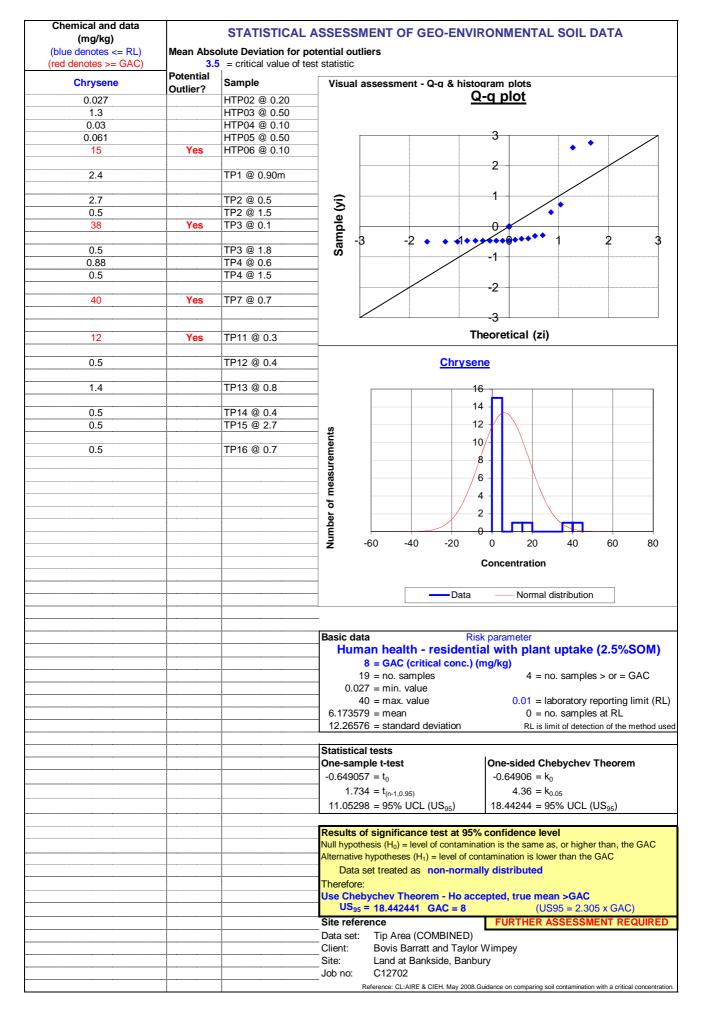






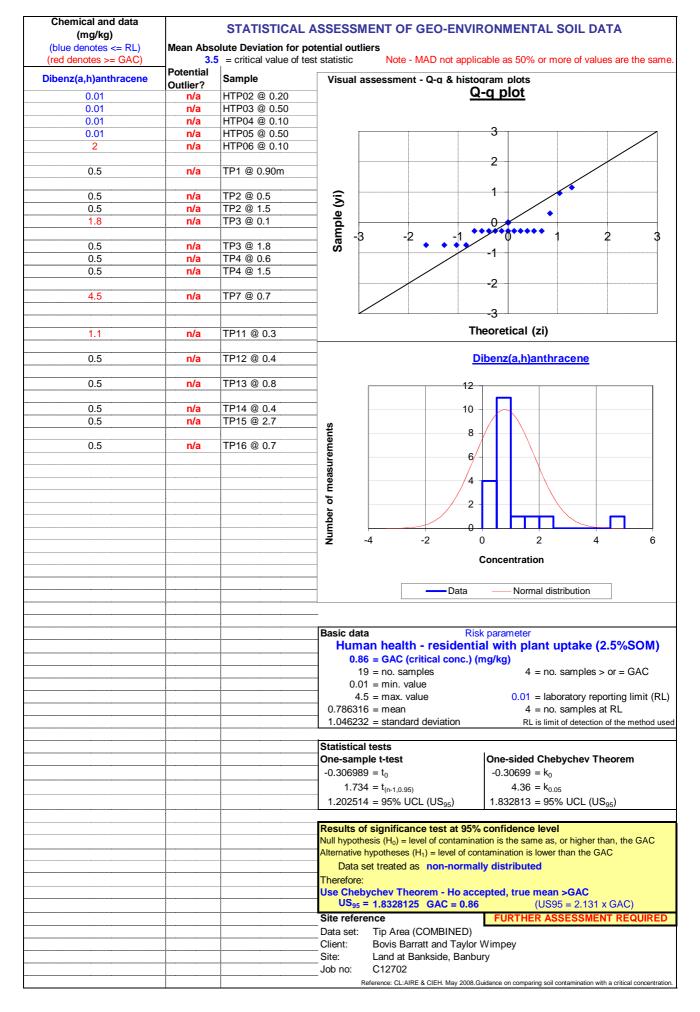


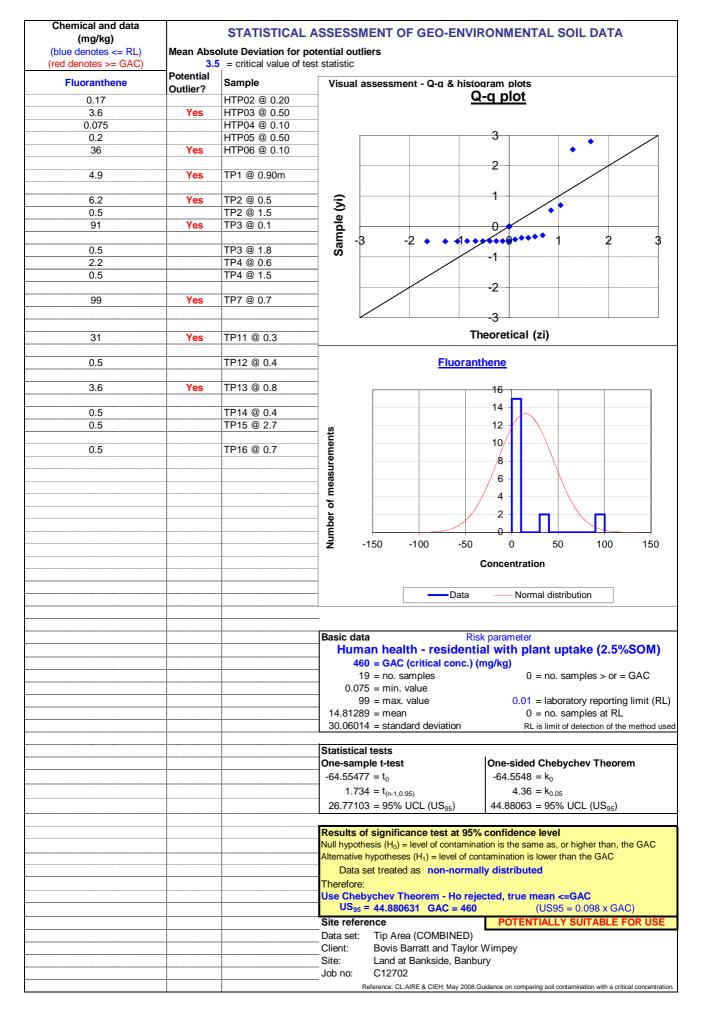








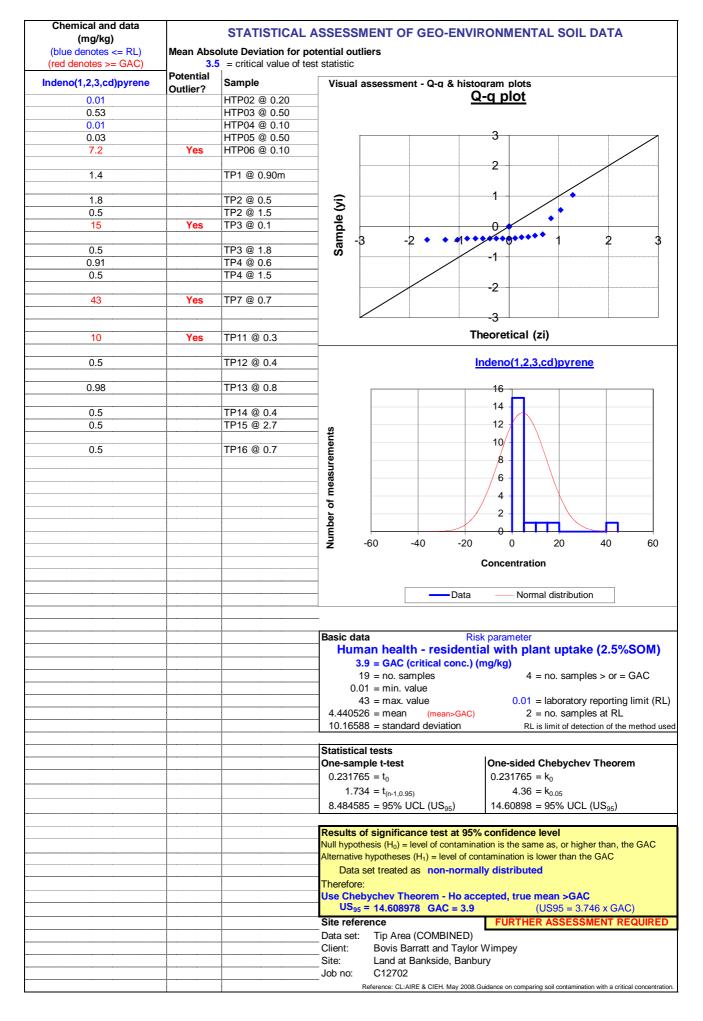




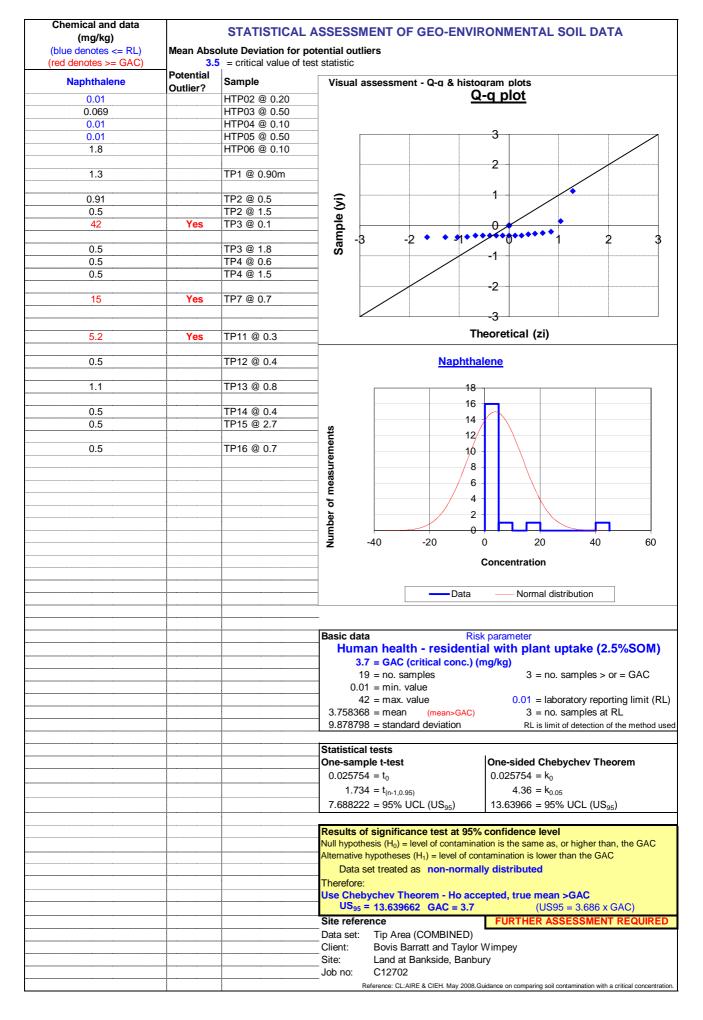




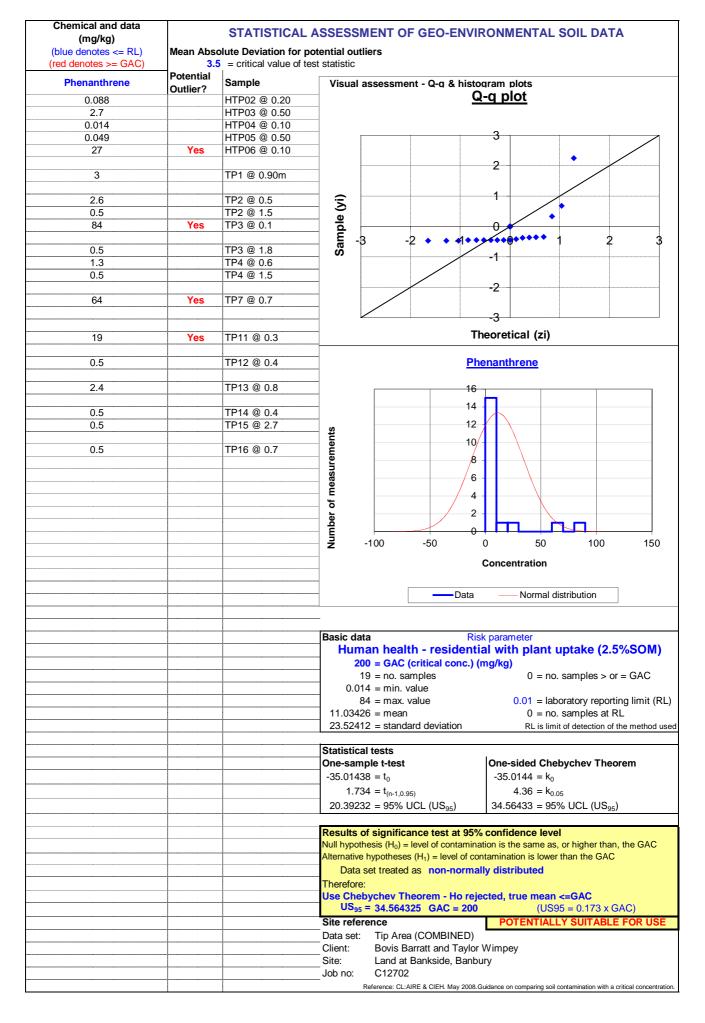
Chemical and data (mg/kg) (blue denotes <= RL) (red denotes >= GAC)		STATISTICAL olute Deviation for p = critical value of to	
(red denotes >= GAC)	Potential		test statistic Note - MAD not applicable as 50% or more of values are the same.
Fluorene	Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.01	n/a	HTP02 @ 0.20	Q-q plot
			<u></u>
0.16	n/a	HTP03 @ 0.50	
0.01	n/a	HTP04 @ 0.10	
0.01	n/a	HTP05 @ 0.50	
2.3	n/a	HTP06 @ 0.10	2
0.5	n/a	TP1 @ 0.90m	
0.5	n/a	TP2 @ 0.5	
0.5	n/a	TP2 @ 1.5	
18	n/a	TP3 @ 0.1	Sample (i) -3 -2 • • • • • • • • • 1 2 3
0.5	n/a	TP3 @ 1.8	S I I I I I I I I I I I I I I I I I I I
0.5	n/a	TP4 @ 0.6	
0.5	n/a	TP4 @ 1.5	-2
9.1	n/a	TP7 @ 0.7	
1.3	n/a	TP11 @ 0.3	Theoretical (zi)
0.5	n/a	TP12 @ 0.4	Fluorene
0.5	n/a	TP13 @ 0.8	
0.5	n/a	TP14 @ 0.4	
0.5	n/a	TP15 @ 2.7	14 - 12
0.5	n/a	TP16 @ 0.7	stemen stemen
			ヹ -20 -10 0 10 20 30 Concentration
			Data — Normal distribution
			Data Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			380 = GAC (critical conc.) (mg/kg)
			19 = no. samples 0 = no. samples > or = GAC
			0.01 = min. value
			18 = max. value 0.01 = laboratory reporting limit (RL)
			1.915263 = mean 3 = no. samples at RL
			4.389821 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-375.4215 = t_0$ $-375.422 = k_0$
			$1.734 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			3.661564 = 95% UCL (US ₉₅) 6.306193 = 95% UCL (US ₉₅)
			Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H_1) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			US ₉₅ = 6.3061928 GAC = 380 (US95 = 0.017 x GAC) Site reference POTENTIALLY SUITABLE FOR USE
			Data set: Tip Area (COMBINED) Client: Bovis Barratt and Taylor Wimpey
			Site: Land at Bankside, Banbury
			Job no: C12702
L	1		Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.



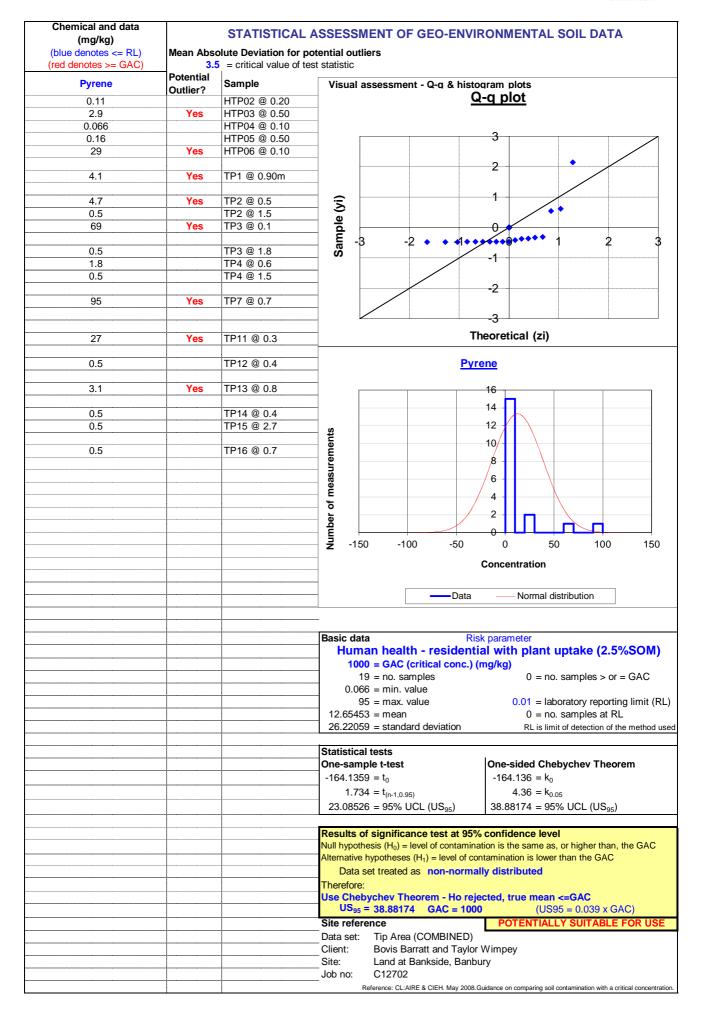




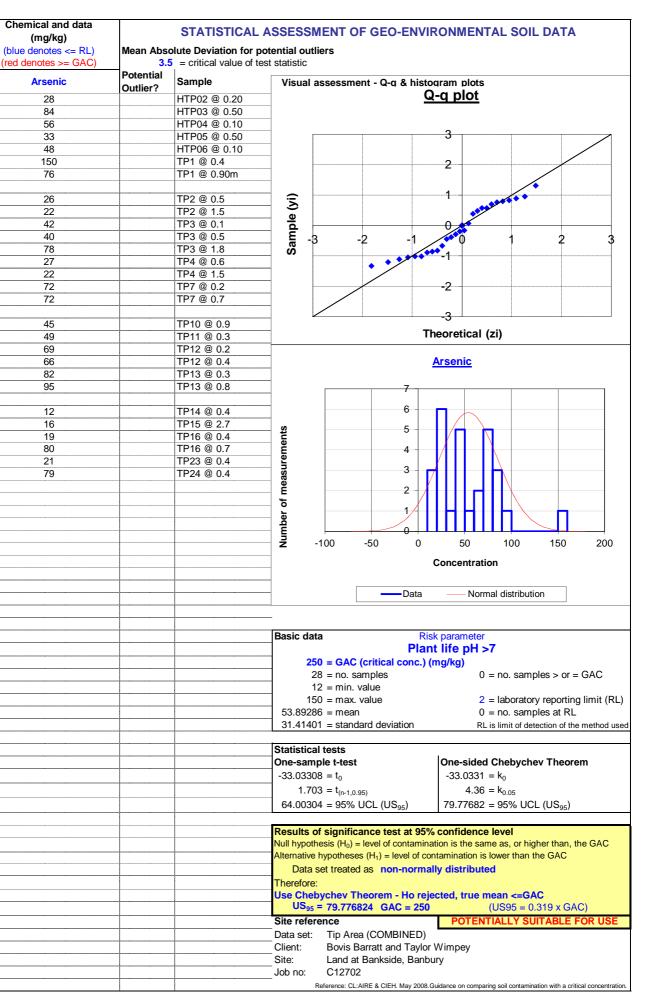




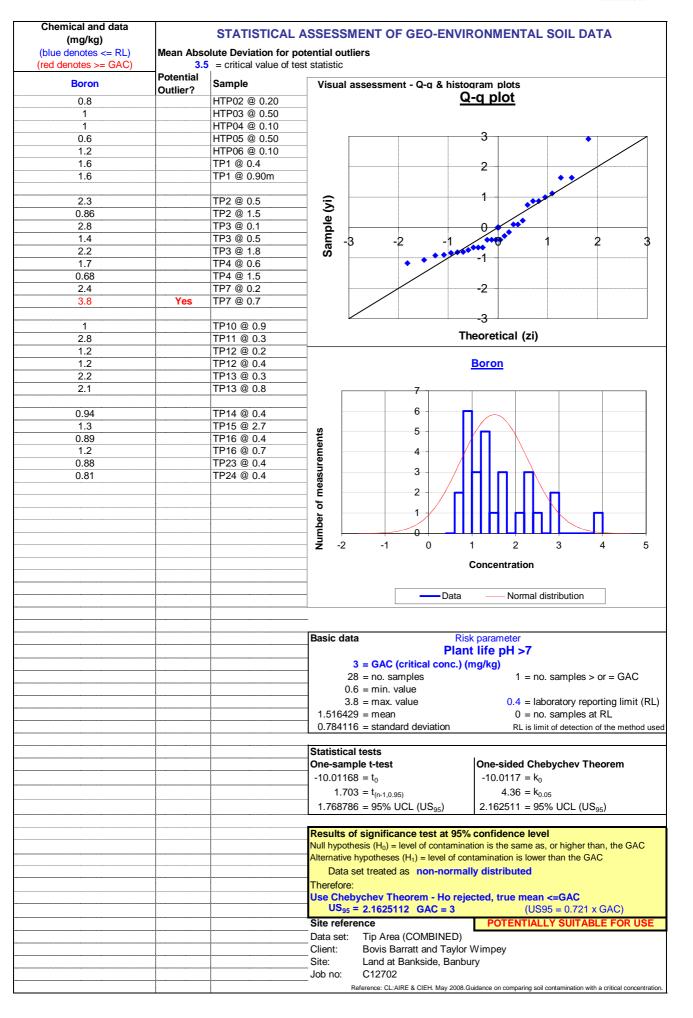






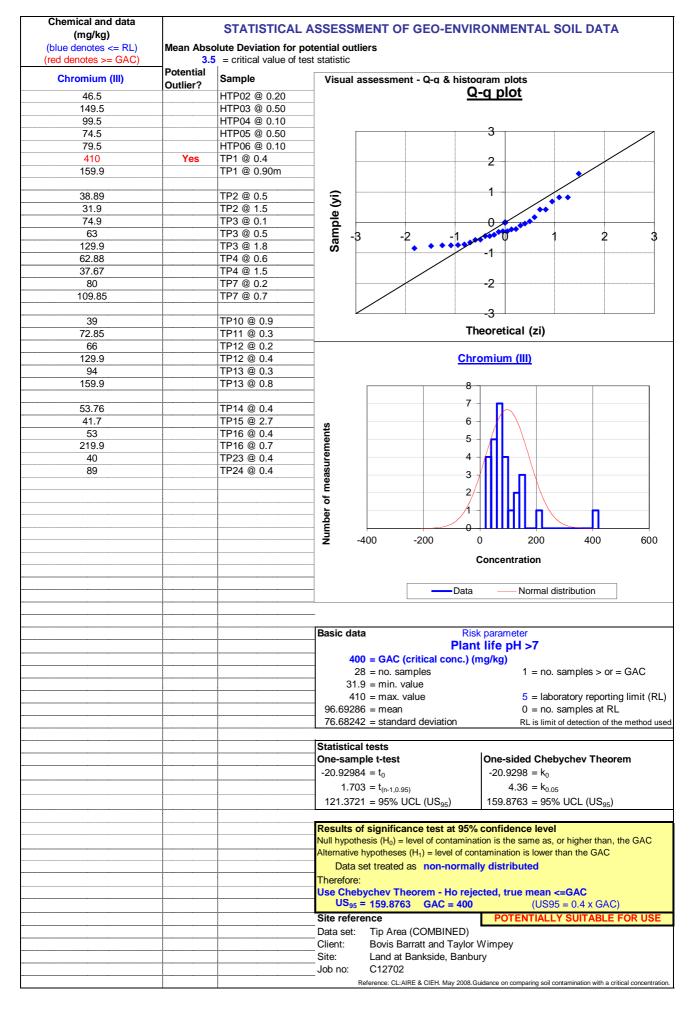


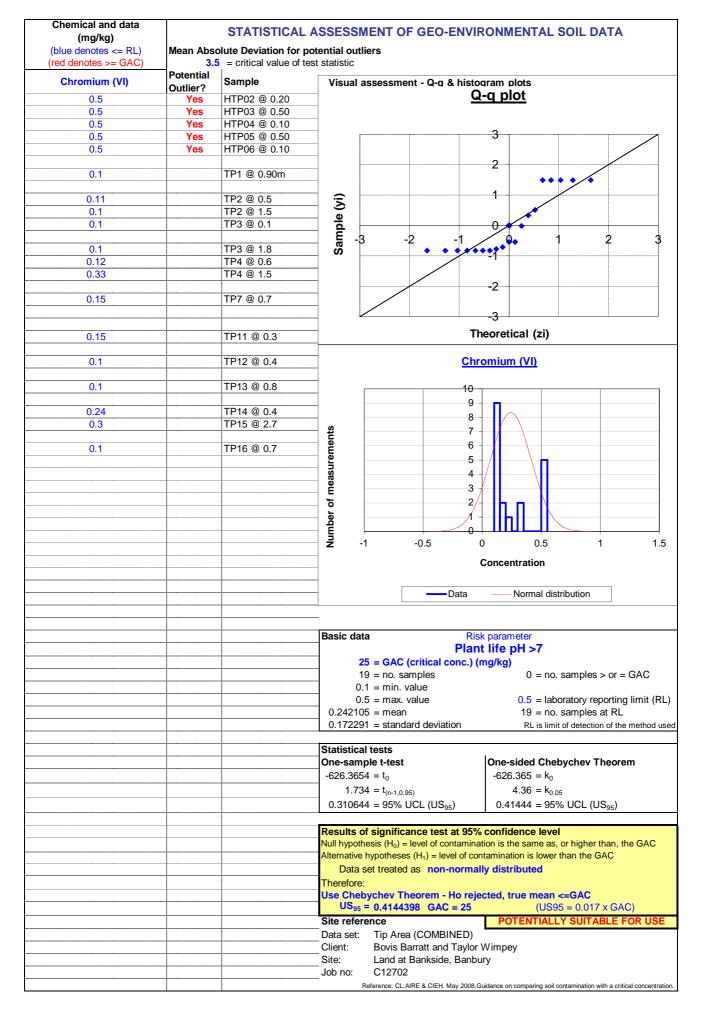




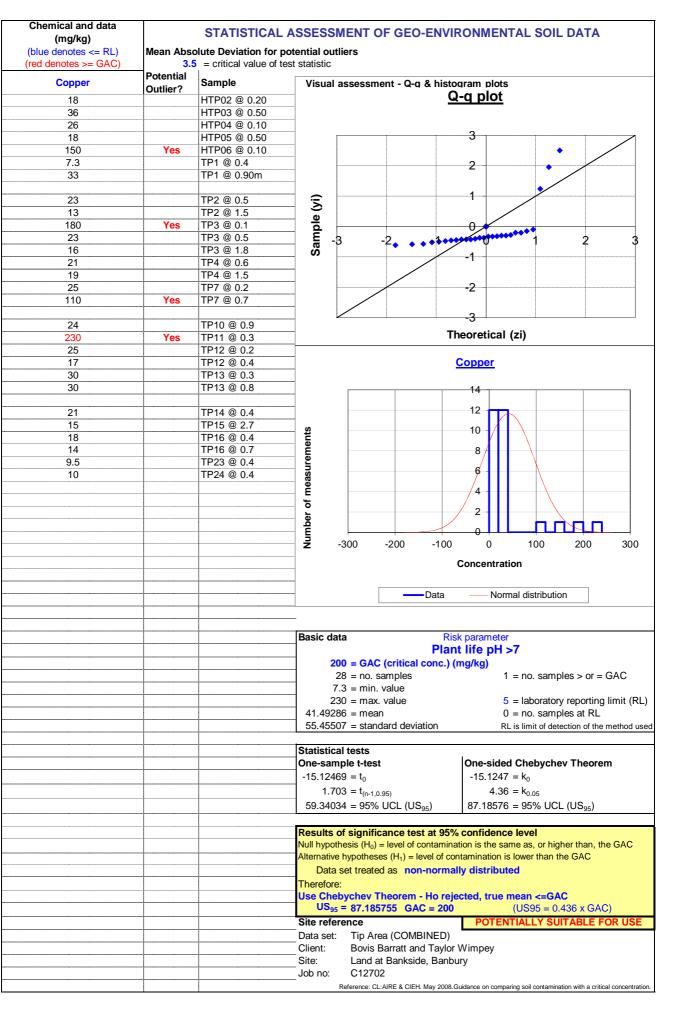




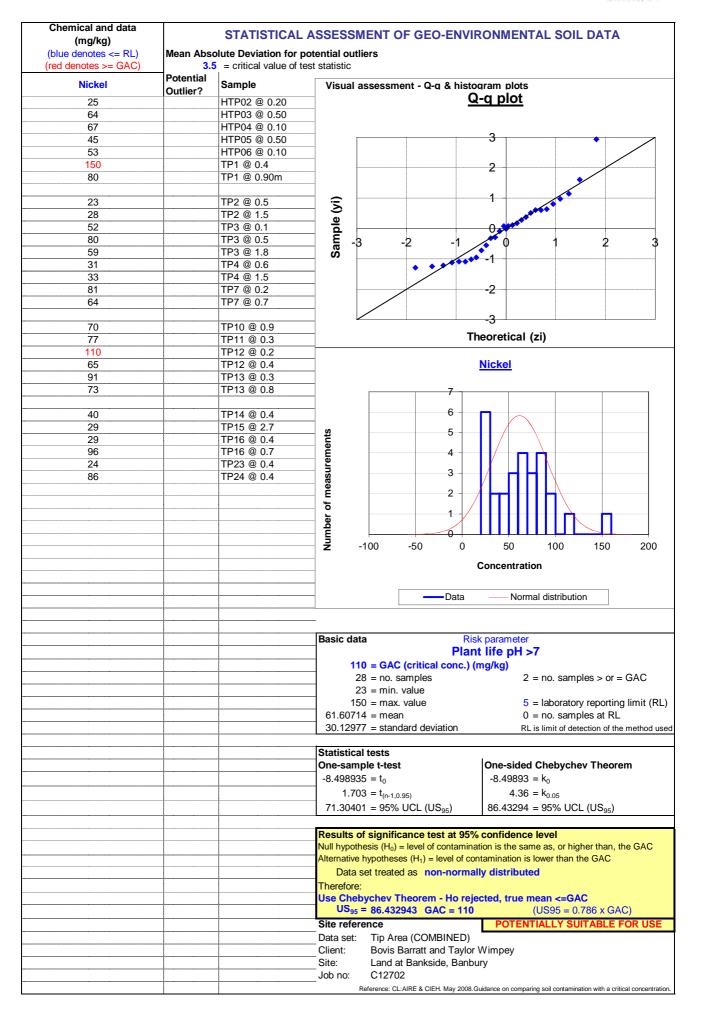












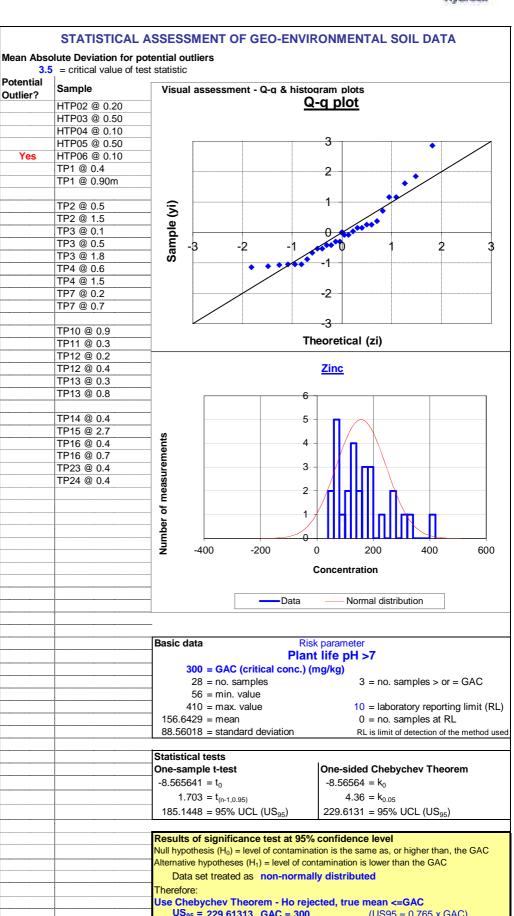


Chemical and data

(mg/kg) (blue denotes <= RL)

(red denotes >= GAC)

Zinc



		3 • • • • • •
Alterna	ative hypotheses $(H_1) =$ level of con	tamination is lower than the GAC
D	ata set treated as non-normal	ly distributed
There	fore:	
Use C	hebychev Theorem - Ho reje	cted, true mean <=GAC
U	S ₉₅ = 229.61313 GAC = 300	(US95 = 0.765 x GAC)
Site r	eference	POTENTIALLY SUITABLE FOR USE
Data	set: Tip Area (COMBINED)	
Client	Bovis Barratt and Taylor	Wimpey
Site:	Land at Bankside, Banbu	ry
Job n	D: C12702	
	Reference: CL:AIRE & CIEH. May 2008.Gu	uidance on comparing soil contamination with a critical concentration.



Assessment of Chemicals of Potential Concern to Human Health

						Soil Type	NAT	NAT	NAT	NAT	NAT	NAT		 		
	All values i	n mg/kg unle	ss otherwis	e stated	Locatio	on & Depth	HTP07	HTP09	HTP10	HTP13	TP19	TP21				
hemical of Potential oncern	Lab. RL	No. Samples	Min. Value	Max. Value	No. Samples > or = GAC	GAC	0.50	0.20	0.50	0.20	0.5	0.4				
rsenic	2	6	10	22	0	32	18	22	18	17	10	21				
eryllium	1	4	1	1	0	51	1	1	1	1						
oron	0.4	6	0.4	0.77	0	290	0.5	0.4	0.5	0.7	0.65	0.77				
admium	0.1	6	0.1	0.5	0	11	0.1	0.1	0.1	0.1	0.5	0.5				
hromium (III)	5	6	38.5	48.5	0	630	48.5	40.5	41.5	38.5	42	45				
hromium (VI)	0.5	4	0.5	0.5	0	4.3	0.5	0.5	0.5	0.5						
opper	5	6	5.9	15	0	2300	15	12	10	12	11	5.9				
ead	5	6	14	29	0	450	26	16	24	29	14	17	 	 		
lercury, inorganic	0.1	6	0.1	0.2	0	170	0.1	0.1	0.1	0.1	0.2	0.2				
lickel	5	6	21	33	0	130	33	22	22	21	31	26				
elenium	0.2	6	0.2	0.3	0	350	0.2	0.2	0.2	0.2	0.3	0.3			1	
anadium	5	4	53	61	0	74	61	58	58	53					1	1
inc	10	6	59	85	0	3700	85	59	66	61	60	69	 			
yanide (free)	0.5	6	0.5	2	0	750	0.5	0.5	0.5	0.5	2	2	 			
henol (total)	0.3	6	0.3	0.5	0	290	0.3	0.3	0.3	0.3	0.5	0.5	 	 		
cenaphthene	0.01	4	0.01	0.01	0	480	0.01	0.01	0.01	0.01			 			
cenaphthylene	0.01	4	0.01	0.01	0	400	0.01	0.01	0.01	0.01			 	 	1	
nthracene	0.01	4	0.01	0.01	0	4900	0.01	0.01	0.01	0.01			 		-	
enz(a)anthracene	0.01	4	0.01	0.023	0	4.7	0.023	0.01	0.012	0.021			 	 		-
enzo(a)pyrene	0.01	4	0.01	0.01	0	0.94	0.01	0.01	0.01	0.01			 	 		
enzo(b)fluoranthene	0.01	4	0.01	0.01	0	6.5	0.01	0.01	0.01	0.01			 	 		
enzo(ghi)perylene	0.01	4	0.01	0.01	0	46	0.01	0.01	0.01	0.01			 	 -		
enzo(k)fluoranthene	0.01	4	0.01	0.01	0	9.6	0.01	0.01	0.01	0.01			 	 		
hrysene	0.01	4	0.01	0.013	0	8	0.013	0.01	0.01	0.012			 	 	1	
ibenz(a,h)anthracene	0.01	4	0.01	0.01	0	0.86	0.01	0.01	0.01	0.01				 		
uoranthene	0.01	4	0.01	0.088	0	460	0.047	0.01	0.031	0.088			 	-		-
uorene	0.01	4	0.01	0.01	0	380	0.01	0.01	0.01	0.01				 -		-
deno(1,2,3,cd)pyrene	0.01	4	0.01	0.01	0	3.9	0.01	0.01	0.01	0.01			 	 		
aphthalene	0.01	4	0.01	0.01	0	3.7	0.01	0.01	0.01	0.01			 	 	+	
henanthrene	0.01	4	0.01	0.018	0	200	0.018	0.01	0.01	0.014						
/rene	0.01	4	0.01	0.061	0	1000	0.035	0.01	0.017	0.061			 			
	Mean	· · ·			Ŭ		0.000						 	 	+	
DC (dimensionless)	0.013						0.023	0.0046	0.019	0.018	0.003081	0.00814	 	 		
OM (calculated)	2.18%						3.97%	0.79%	3.28%	3.10%	0.53%	1.40%		+	+	
H (su)	7.3						7.1	6.9	7.3	7.3	7.6	7.5	 	 	+	
1 (34)	1.5	l	L	I	L		1.1	1 0.3	1.5	1.5	1.0	1.5	 	 	l	

Client: Bovis Barratt and Taylor Wimpy Site: Land at Bankside, Banbury Job no: C12702 considered as being at the detection limit for the purposes of statistical analysis, as a conservative estimate. Values in red are equal to, or greater than, the generic assessment criterion (GAC). MG denotes Made Ground NAT denotes natural ground Hydrock

Assessment of Chemicals of Potential Concern to Plant Life

						Soil Type	NAT	NAT	NAT	NAT	NAT	NAT		
	All values i	Locati	on & Depth	HTP07	HTP09	HTP10	HTP13	TP19	TP21					
Chemical of Potential Concern	Lab. RL	No. Samples	Min. Value	Max. Value	No. Samples > or = GAC	GAC	0.50	0.20	0.50	0.20	0.5	0.4		
Arsenic	2	6	10	22	0	250	18	22	18	17	10	21		 -
Boron	0.4	6	0.4	0.77	0	3	0.5	0.4	0.5	0.7	0.65	0.77		
Chromium (III)	5	6	38.5	48.5	0	400	48.5	40.5	41.5	38.5	42	45		
Chromium (VI)	0.5	4	0.5	0.5	0	25	0.5	0.5	0.5	0.5				
Copper	5	6	5.9	15	0	135	15	12	10	12	11	5.9		
Nickel	5	6	21	33	0	75	33	22	22	21	31	26		1
Zinc	10	6	59	85	0	300	85	59	66	61	60	69		
	Mean													
pH (su)	7.3			1	1		7.1	6.9	7.3	7.3	7.6	7.5	1	 1

Data set: CMF & DF Client: Bovis Barratt and Taylor Wimpy Site: Land at Bankside, Banbury Job no: C12702

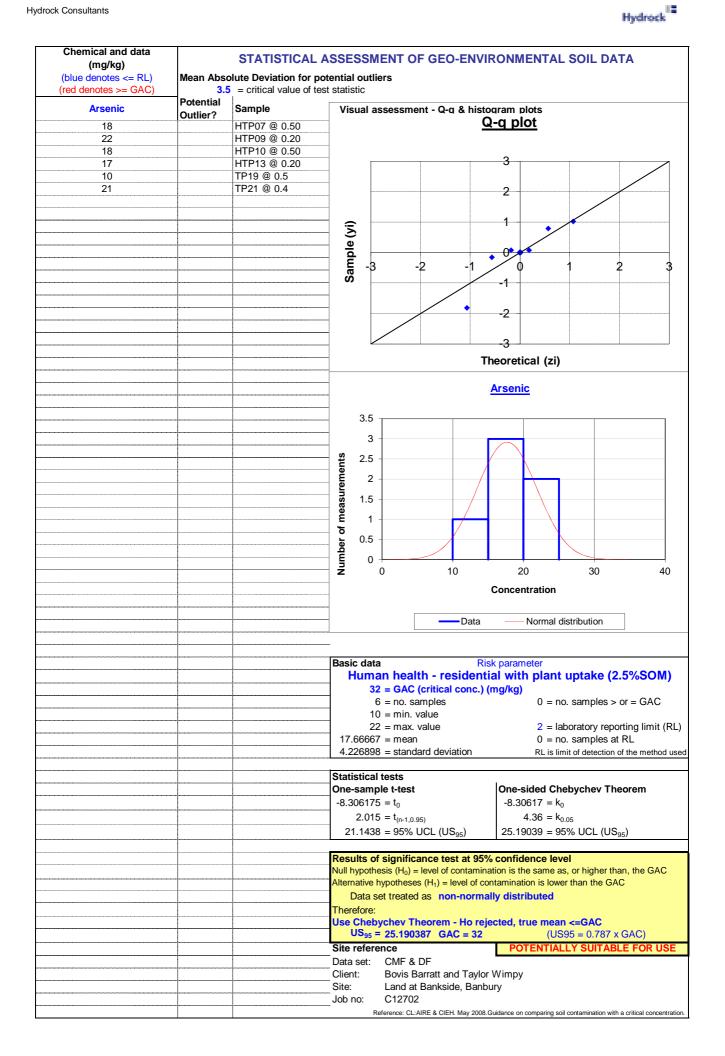
 Legend:
 Values in blue are at or below the laboratory reporting limit (where a single value is indicated) and are considered as being at the detection limit for the purposes of statistical analysis, as a conservative estimate.

 Values in red are equal to, or greater than, the generic assessment criterion (GAC).

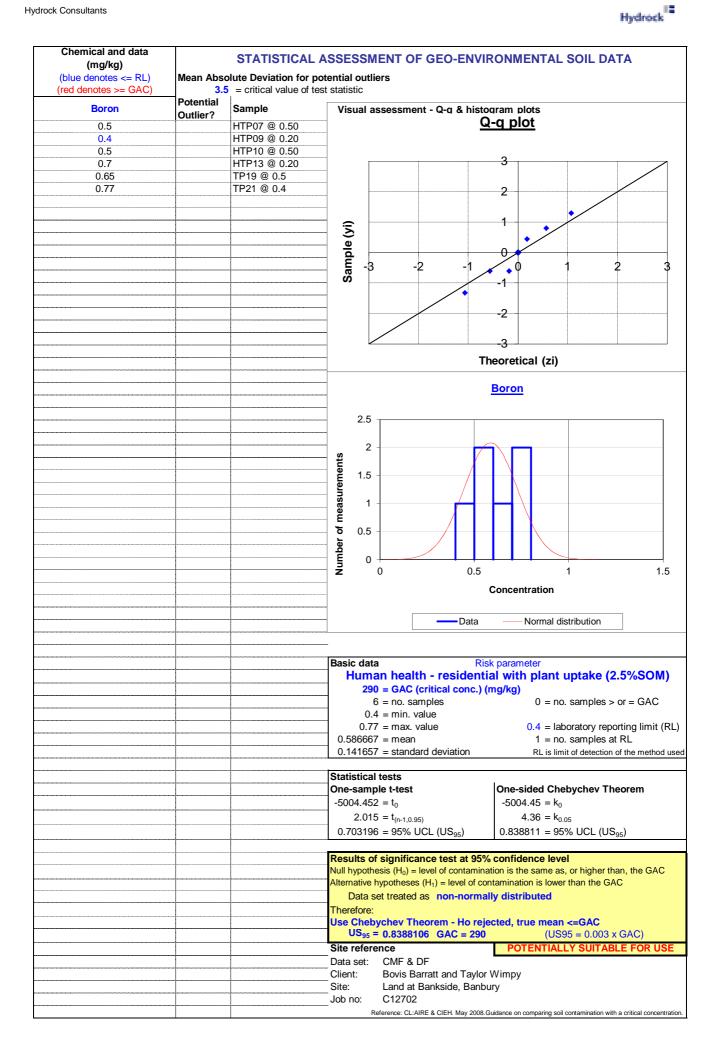
 MG denotes Made Ground

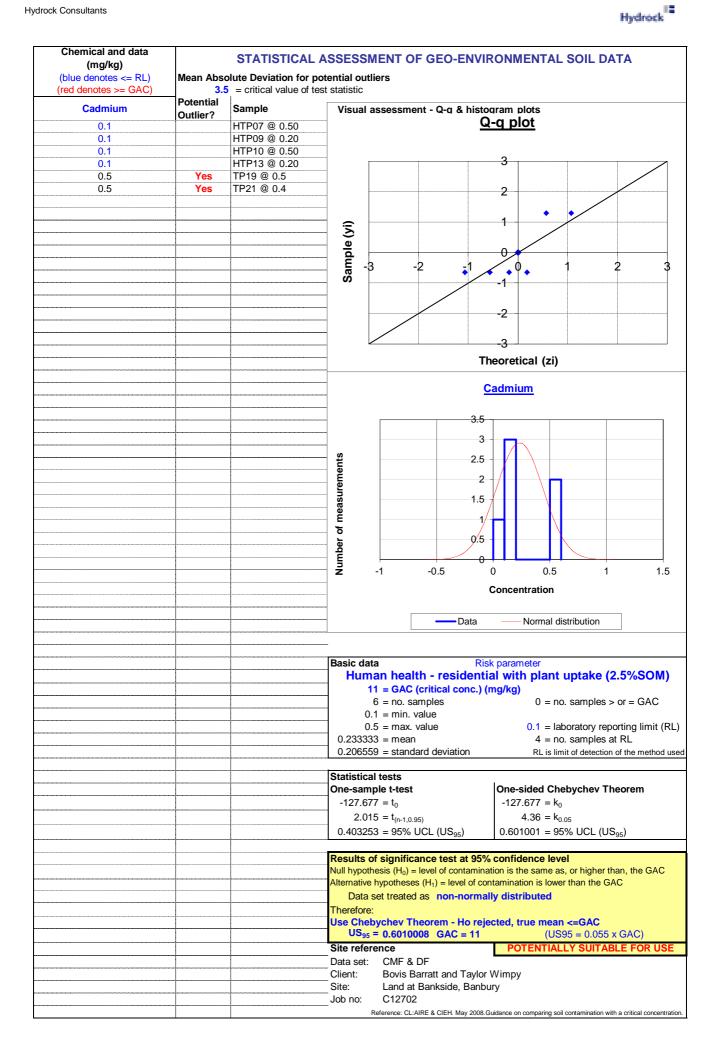
 NAT denotes natural ground

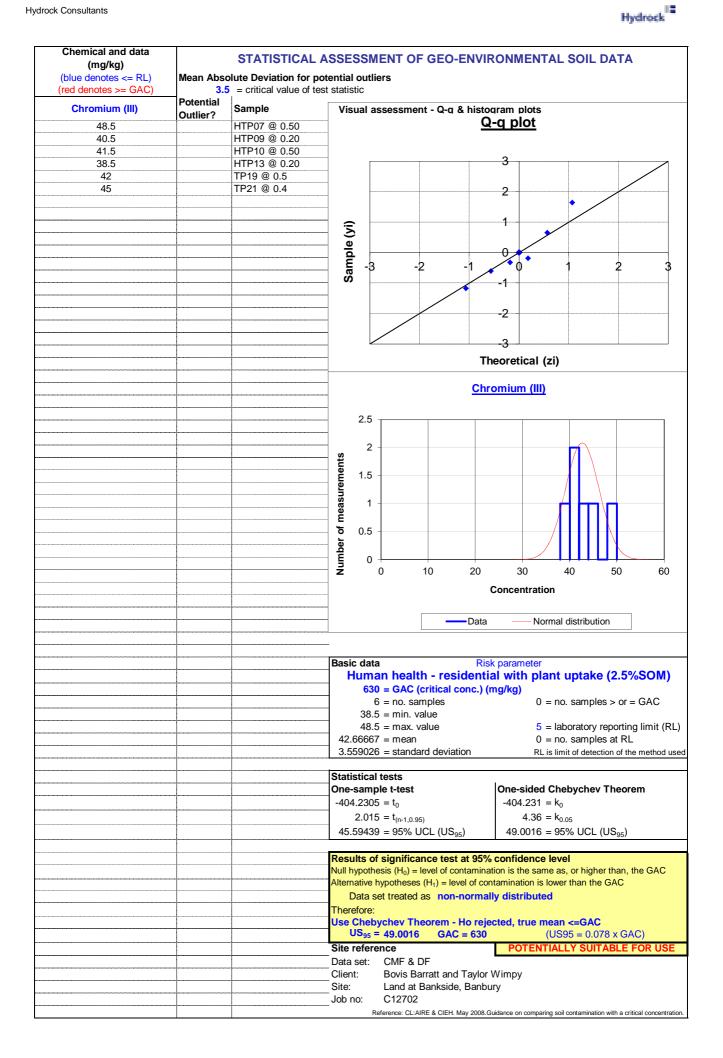
Hydrock



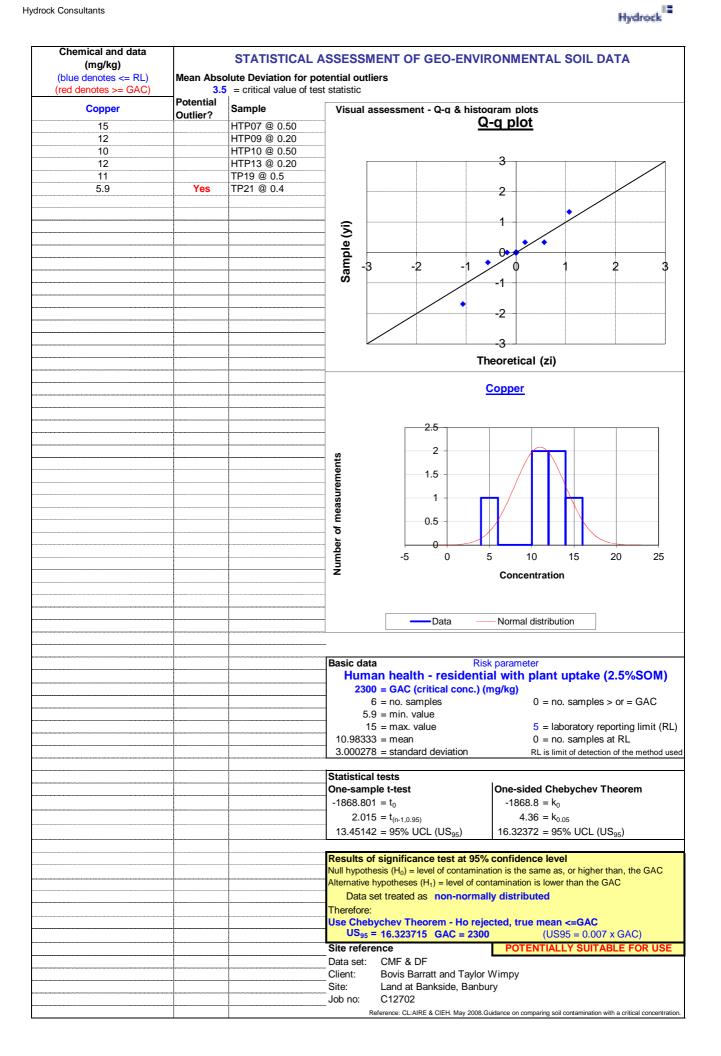
Chemical and data (mg/kg)		STATISTICAL	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL)		plute Deviation for p	
(red denotes >= GAC) Beryllium	3.8 Potential	<pre>= critical value of te Sample</pre>	est statistic Note - MAD not applicable as 50% or more of values are the same. Visual assessment - Q-g & histogram plots
1	Outlier? n/a	HTP07 @ 0.50	
1	n/a	HTP09 @ 0.20	
1	n/a	HTP10 @ 0.50	
1	n/a	HTP13 @ 0.20	3
		-	
			- Š Š
			Sample (ji)
			-
			-2
			-3
			Theoretical (zi)
			Demillion
			Beryllium
			3.5
			2.5 2 1.5 0.5 0 1 1 1 1 1 1 1 1 1 1 1 1
			Concentration
			Data Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			51 = GAC (critical conc.) (mg/kg) 4 = no. samples 0 = no. samples > or = GAC
			1 = min. value
			1 = max. value 1 = laboratory reporting limit (RL)
			1 = mean 4 = no. samples at RL 5E-11 = standard deviation RL is limit of detection of the method used
			Statistical tests One-sample t-test One-sided Chebychev Theorem
	+		One-sample t-test One-sided Chebychev Theorem -2E+12 = t ₀ -2E+12 = k ₀
	-	1	$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			1 = 95% UCL (US ₉₅) $1 = 95%$ UCL (US ₉₅)
			Results of significance test at 95% confidence level Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses (H ₁) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			$US_{95} = 1 \qquad GAC = 51 \qquad (US95 = 0.02 \times GAC)$
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy Site: Land at Bankside, Banbury
			Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.





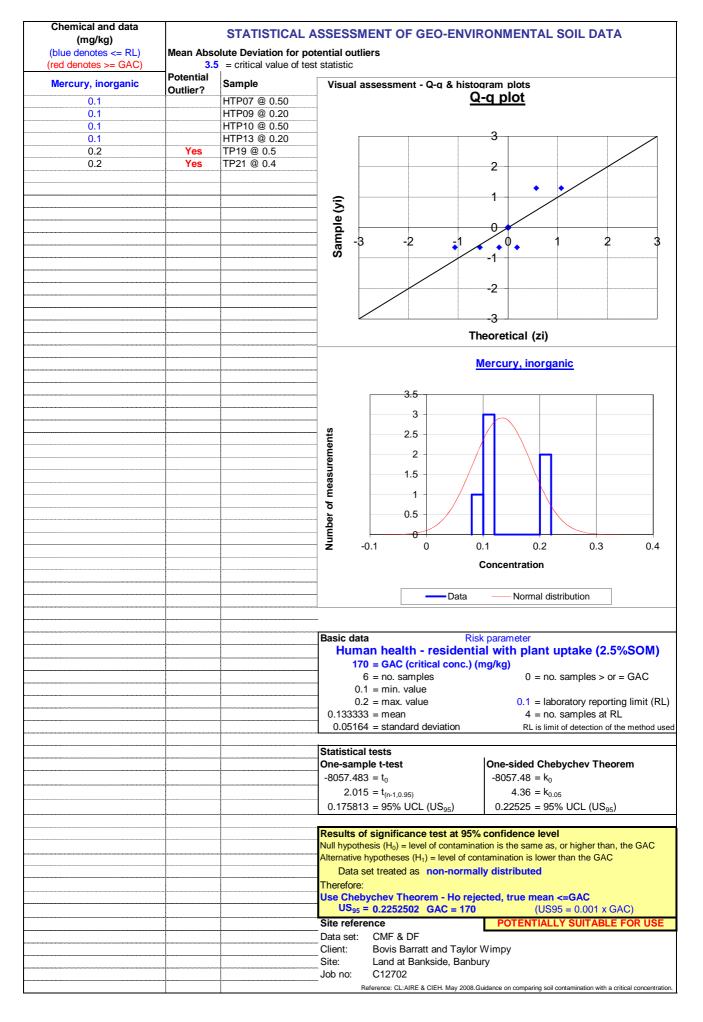


Chemical and data (mg/kg)		STATISTICAL	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL) (red denotes >= GAC)		blute Deviation for po = critical value of te	
Chromium (VI)	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.5	n/a	HTP07 @ 0.50	<u>Q-q plot</u>
0.5	n/a	HTP09 @ 0.20	
0.5	n/a n/a	HTP10 @ 0.50 HTP13 @ 0.20	3
0.0	11/4	1117 13 @ 0.20	
			2
			Sample (y)
			\neg
			-2
			Theoretical (zi)
			Chromium ()/)
			Chromium (VI)
			3.5
			- 월 2.5
			5 1
			2.5 2 1.5 0 0 0,5 0,5 0,5 0,5 0,5 0,5 0,5 0,5 0,
			Concentration
			Data Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			4 = no. samples 0 = no. samples > or = GAC
			0.5 = min. value
			0.5 = max. value 0.5 = laboratory reporting limit (RL)
			0.5 = mean 4 = no. samples at RL 5E-12 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem -1.52E+12 = t ₀ -1.5E+12 = k ₀
			$\begin{array}{c} -1.52\pm12 = t_0 \\ 2.353 = t_{(n-1,0.95)} \end{array} \qquad \begin{array}{c} -1.5\pm12 = k_0 \\ 4.36 = k_{0.05} \end{array}$
L			$0.5 = 95\% \text{ UCL } (\text{US}_{95}) \qquad 0.5 = 95\% \text{ UCL } (\text{US}_{95})$
			Results of significance test at 95% confidence level Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses (H_0) = level of contamination is the same as, of higher than, the GAC Alternative hypotheses (H_1) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC US ₉₅ = 0.5 GAC = 4.3 (US95 = 0.116 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy Site: Land at Bankside, Banbury
			Site: Land at Bankside, Banbury Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008.Guidance on comparing soil contamination with a critical concentration.

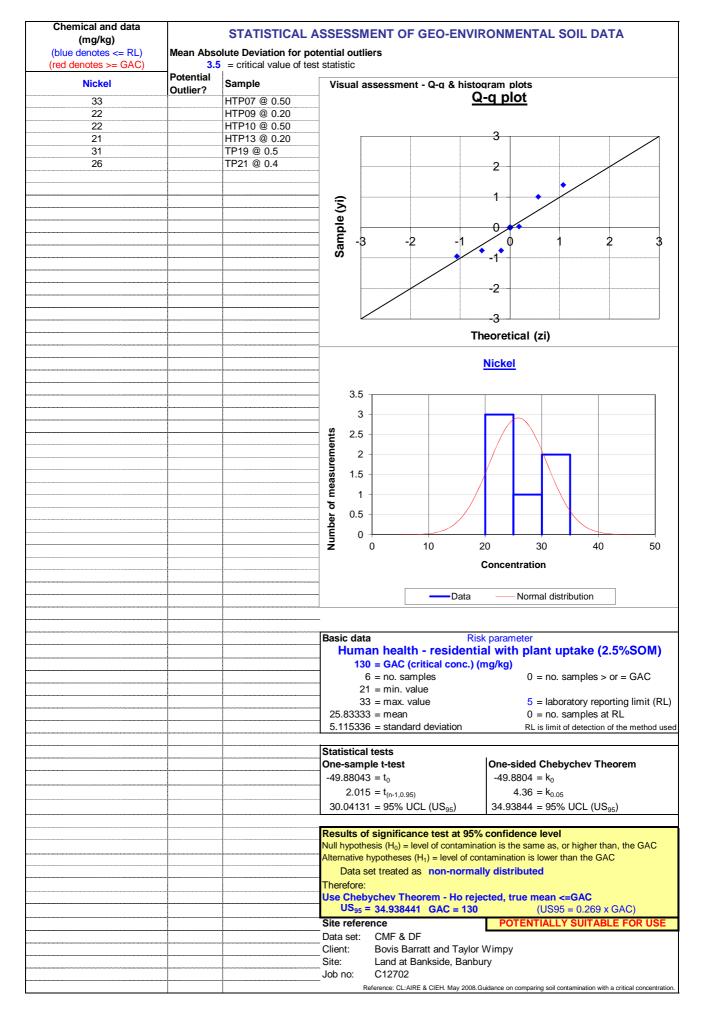


Chemical and data (mg/kg)		STATISTICAL A	SSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL)		olute Deviation for po	
(red denotes >= GAC)		= critical value of tes	t statistic
Lead	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots
26	Outlier	HTP07 @ 0.50	<u>Q-q plot</u>
16		HTP09 @ 0.20	
24		HTP10 @ 0.50	
29		HTP13 @ 0.20	3
14		TP19 @ 0.5	
17		TP21 @ 0.4	
		1121 8 0.1	2
			Sample (j)
			-1
			-2
			-3
			Theoretical (zi)
			Lead
			2.5
			2.3
			1.5 1 .5
			5 5
			-10 0 10 20 30 40 50
			Z Concentration
			Concentration
			-
			~
			Data Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			450 = GAC (critical conc.) (mg/kg)
			6 = no. samples 0 = no. samples > or = GAC
			14 = min. value
			29 = max. value 5 = laboratory reporting limit (RL) 21 = mean 0 = no. samples at RL
			21 = mean 0 = no. samples at RL 6.131884 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-171.3717 = t_0$ $-171.372 = k_0$
			$2.015 = t_{(n-1,0.95)}$ $4.36 = k_{0.05}$
	-		26.04421 = 95% UCL (US ₉₅) $31.91452 = 95%$ UCL (US ₉₅)
			Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses (H_1) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			US ₉₅ = 31.914524 GAC = 450 (US95 = 0.071 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			_Job no: C12702
	1		Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.

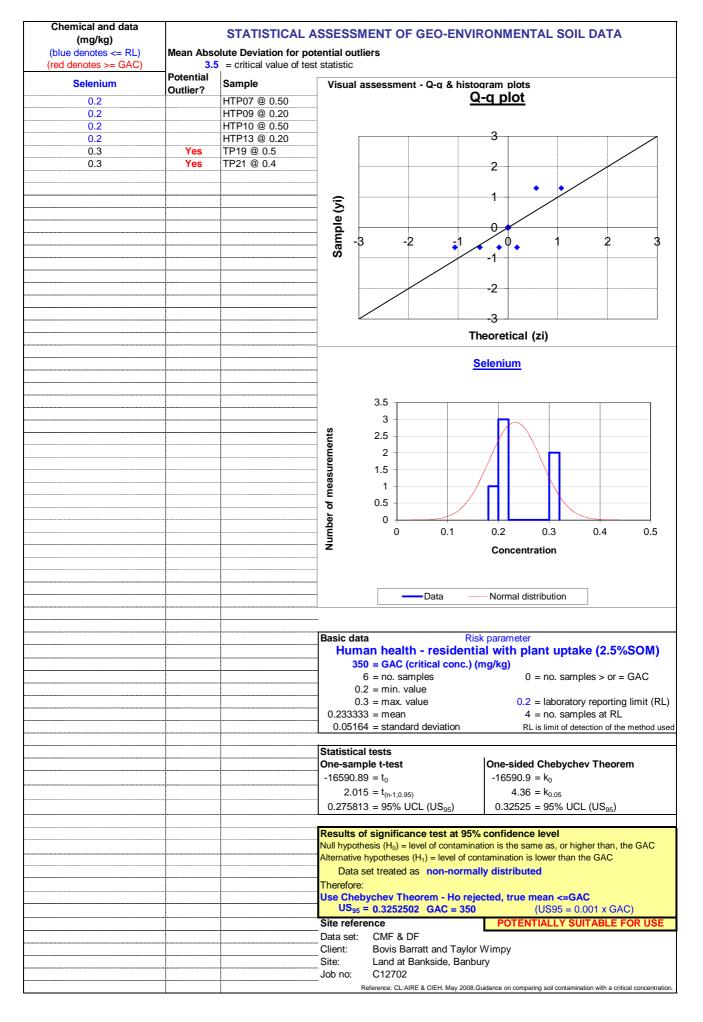








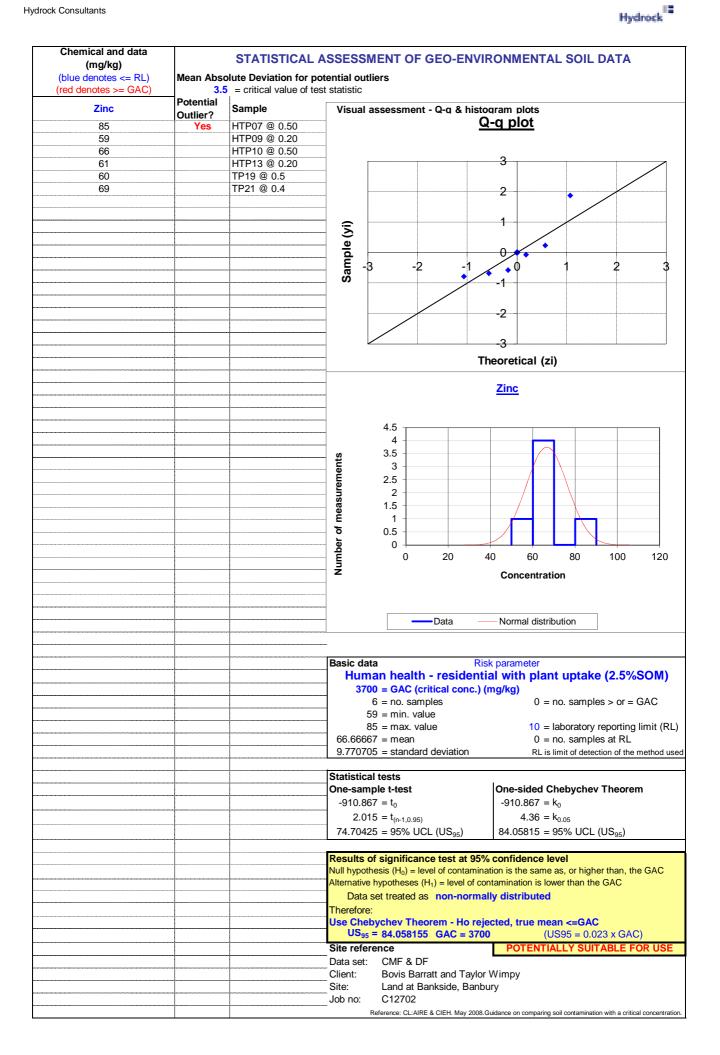


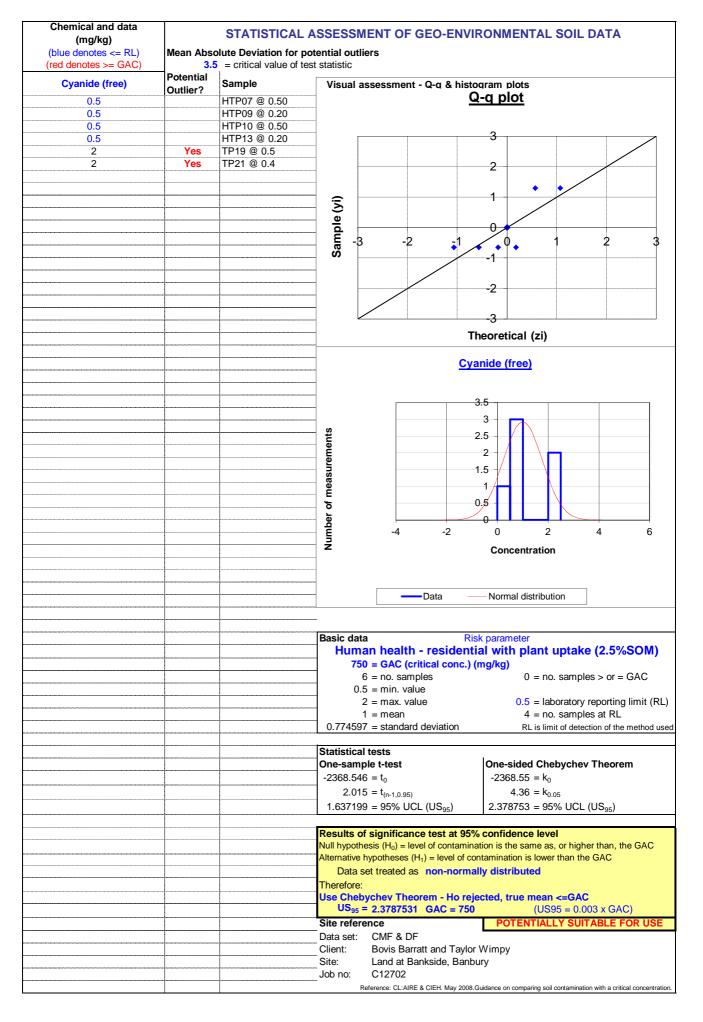




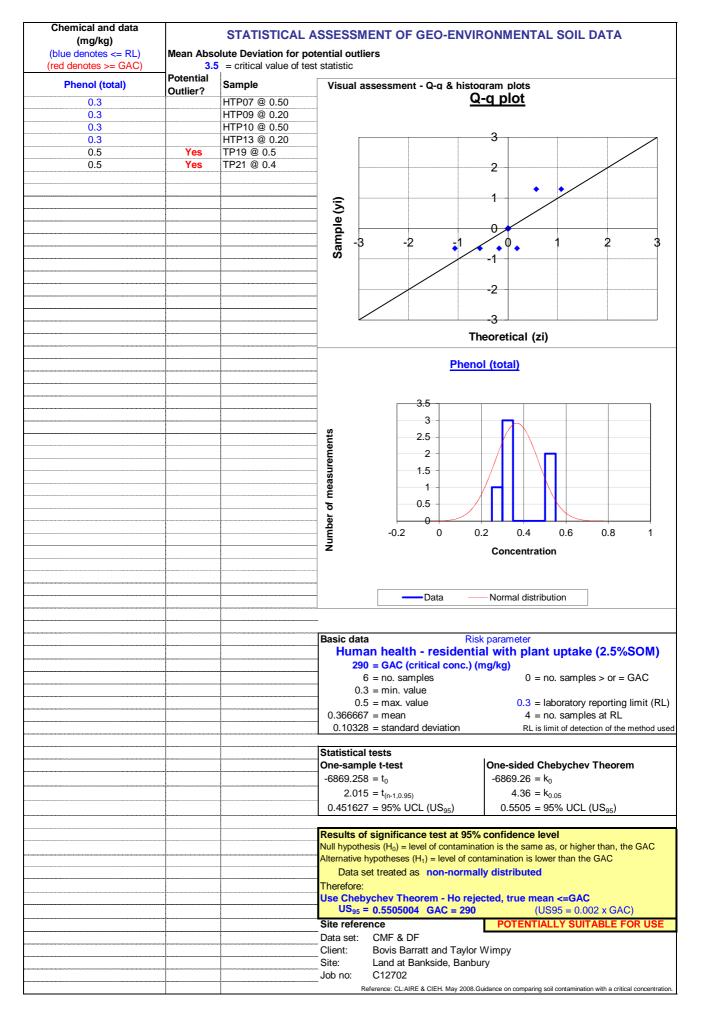
Chemical and data (mg/kg)		STATISTICAL	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL) (red denotes >= GAC)		blute Deviation for p = critical value of te	
Vanadium	Potential	Sample	Visual assessment - Q-q & histogram plots
61	Outlier?	HTP07 @ 0.50	<u>Q-q plot</u>
58 58		HTP09 @ 0.20 HTP10 @ 0.50	
53		HTP10 @ 0.50 HTP13 @ 0.20	3
			2
			Bandhar San
			· · · · · · · · · · · · · · · · · · ·
			-2
			Theoretical (zi)
			Vanadium
			2.5
			st
			Z Concentration
			Data Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			74 = GAC (critical conc.) (mg/kg) 4 = no. samples 0 = no. samples > or = GAC
			53 = min. value
			61 = max. value 5 = laboratory reporting limit (RL) 57.5 = mean 0 = no. samples at RL
			57.5 = mean 0 = no. samples at RL 3.316625 = standard deviation RL is limit of detection of the method used
			Statistical tests One-sample t-test One-sided Chebychev Theorem
			$-9.949874 = t_0 \qquad -9.94987 = k_0$
			$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			$61.40201 = 95\% \text{ UCL } (\text{US}_{95}) \qquad 64.73024 = 95\% \text{ UCL } (\text{US}_{95})$
			Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses (H ₁) = level of contamination is lower than the GAC Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC US ₉₅ = 64.730242 GAC = 74 (US95 = 0.875 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy Site: Land at Bankside, Banbury
			Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.













Chemical and data (mg/kg)	STATISTICAL ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA			
(blue denotes <= RL)	Mean Absolute Deviation for potential outliers 3.5 = critical value of test statistic Note - MAD not applicable as 50% or more of values are the s			
(red denotes >= GAC) Acenaphthene	Potential Outlier?	Sample	Note - MAD not applicable as 50% or more of values are the same. Visual assessment - Q-q & histogram plots	
0.01	n/a	HTP07 @ 0.50	Q-q plot	
0.01	n/a	HTP09 @ 0.20		
0.01	n/a	HTP10 @ 0.50		
0.01	n/a	HTP13 @ 0.20		
0.01	- Tira	1111 13 @ 0.20		
			2	
	-			
			Sample (y)	
			-1	
			-2	

			Theoretical (zi)	
			Acenaphthene	
			3.5	
			3	
			2.5	
			5 0.5	
			0.01 0.01 0.01 0.01 0.01 0.01	
			2 Concentration	
			Data Normal distribution	
			Basic data Risk parameter	
			Human health - residential with plant uptake (2.5%SOM)	
		1	480 = GAC (critical conc.) (mg/kg)	
		+	460 = GAC (childai conc.) (ing/kg) 4 = no. samples $0 = no. samples > or = GAC$	
			4 = 10. samples $0 = 10.$ samples $> 0 = 0.01$	
			0.01 = max. value 0.01 = laboratory reporting limit (RL)	
			0.01 = max. Value $0.01 = nabolatory reporting mint (NL)0.01 = mean$ $4 = no. samples at RL$	
			5E-13 = standard deviation RL is limit of detection of the method used	
		1		
	-		Statistical tests	
			One-sided Chebychev Theorem	
	1		$-1.92E+15 = t_0$ $-1.9E+15 = k_0$	
			$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$	
			$0.01 = 95\% \text{ UCL } (\text{US}_{95}) \qquad 0.01 = 95\% \text{ UCL } (\text{US}_{95})$	
			0.01 = 30% 000 (00000) $0.01 = 30% 000 (00000)$	
			Populta of significance test at 05% confidence level	
			Results of significance test at 95% confidence level Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC	
			Alternative hypotheses (H_0) = level of contamination is lower than the GAC	
			Data set treated as non-normally distributed	
			Therefore:	
			Use Chebychev Theorem - Ho rejected, true mean <=GAC US ₉₅ = 0.01 GAC = 480 (US95 = 0 x GAC)	
			Site reference POTENTIALLY SUITABLE FOR USE	
			Data set: CMF & DF	
			Client: Bovis Barratt and Taylor Wimpy	
			Site: Land at Bankside, Banbury	
			Job no: C12702	
		1	Reference: CL:AIRE & CIEH. May 2008.Guidance on comparing soil contamination with a critical concentration.	

Chemical and data (mg/kg)		STATISTICAL	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL) (red denotes >= GAC)		olute Deviation for p 5 = critical value of to	
Acenaphthylene	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.01	n/a	HTP07 @ 0.50	Q-q plot
0.01	n/a	HTP09 @ 0.20	
0.01	n/a	HTP10 @ 0.50	
0.01	n/a	HTP13 @ 0.20	
0.01		1111 10 0 0.20	
			2
			$-\overline{\mathbf{S}}$
			Beneficial State S
			1 -1
			-2
			.
			Theoretical (zi)
			Acenaphthylene
			3.5
			3
			- \$ 2.5
			2.5 2 1.5 0 0.5 0 0.01 0.01 0.01 0.01 0.01 0.
			2 0.01 0.01 0.01 0.01 0.01 0.01
			Concentration
			·····
			Data Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			400 = GAC (critical conc.) (mg/kg)
			4 = no. samples $0 = no. samples > or = GAC$
			0.01 = min. value
			0.01 = max. value 0.01 = laboratory reporting limit (RL)
			0.01 = mean 4 = no. samples at RL
			5E-13 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-1.6E+15 = k_0$ $-1.6E+15 = k_0$
			$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			0.01 = 95% UCL (US ₉₅) $0.01 = 95%$ UCL (US ₉₅)

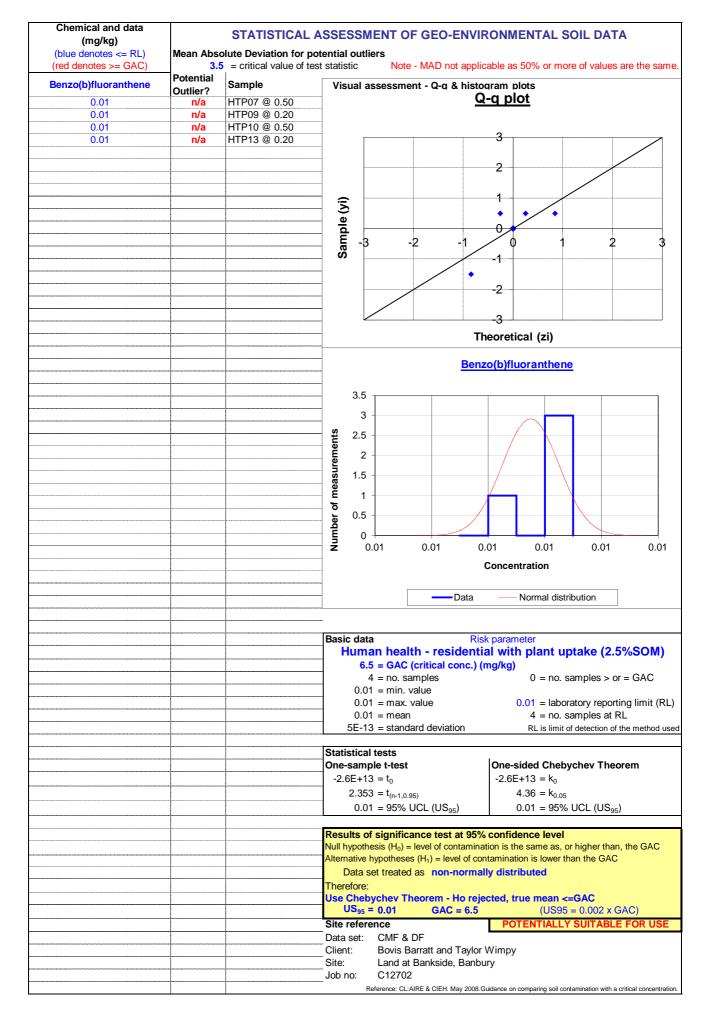
			Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses (H ₁) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			US ₉₅ = 0.01 GAC = 400 (US95 = 0 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008.Guidance on comparing soil contamination with a critical concentration.
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Chemical and data (mg/kg)		STATISTICAL	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL) (red denotes >= GAC)		olute Deviation for po = critical value of te	
Anthracene	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.01	n/a	HTP07 @ 0.50	<u>Q-q plot</u>
0.01	n/a	HTP09 @ 0.20	
0.01	n/a	HTP10 @ 0.50	
0.01	n/a	HTP13 @ 0.20	3
			2
			Sample (ji)
			-1
			-2
			Theoretical (zi)
			Anthreesen
			<u>Anthracene</u>
			3.5
			3
			- £ 2.5
			5
			Z 0.01 0.01 0.01 0.01 0.01 0.01
			Concentration
			Concentration
			DataNormal distribution
			-
			Basic data Risk parameter
	Ì		Human health - residential with plant uptake (2.5%SOM)
			4900 = GAC (critical conc.) (mg/kg)
			4 = no. samples 0 = no. samples > or = GAC 0.01 = min. value
			0.01 = mm. value 0.01 = laboratory reporting limit (RL)
			0.01 = mean 4 = no. samples at RL
			5E-13 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-2E+16 = k_0$
			$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			0.01 = 95% UCL (US ₉₅) 0.01 = 95% UCL (US ₉₅)
			Depute of similiances (act of 00% and the set
			Results of significance test at 95% confidence level Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses (H_1) = level of contamination is the same as, of higher than, the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC US ₉₅ = 0.01 GAC = 4900 (US95 = 0 x GAC)
			US ₉₅ = 0.01 GAC = 4900 (US95 = 0 x GAC) Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			_Job no: C12702
1			Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.

Chemical and data (mg/kg)		STATISTICAL	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL)	Mean Abso	olute Deviation for p	potential outliers
(red denotes >= GAC)	3.5	= critical value of te	
Benz(a)anthracene	Potential	Sample	Visual assessment - Q-q & histogram plots
0.023	Outlier?	HTP07 @ 0.50	Q-q plot
0.01		HTP09 @ 0.20	
0.012		HTP10 @ 0.50	
0.021		HTP13 @ 0.20	
			2
			-1 -1
			-2
			3
			Theoretical (zi)
			Benz(a)anthracene
			_
			- w 2
			1.5 1.5 1.5 0.5 0.5 0.01 0.02 0.03 0.04 0.05
			5 0.5
			- Z -0.02 -0.01 0 0.01 0.02 0.03 0.04 0.05
			Concentration
			Data — Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			4.7 = GAC (critical conc.) (mg/kg)
			4 = no. samples 0 = no. samples > or = GAC
			0.01 = min. value 0.023 = max. value 0.01 = laboratory reporting limit (RL)
			0.0165 = mean $1 = no. samples at RL$
			0.006455 = standard deviation RL is limit of detection of the method used
			01-01-01-01
			Statistical tests One-sample t-test One-sided Chebychev Theorem
			$-1451.129 = t_0$ $-1451.13 = k_0$
			$\begin{array}{c} 1401125 - k_0 \\ 2.353 = t_{(n-1,0.95)} \\ \end{array} $
		-	0.024094 = 95% UCL (US ₉₅) $0.030572 = 95%$ UCL (US ₉₅)
			Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses (H_1) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			$US_{95} = 0.0305718$ GAC = 4.7 (US95 = 0.007 x GAC)
	<u> </u>		Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury Job no: C12702
			SOB TIO. CT27 02 Reference: CL:AIRE & CIEH. May 2008.Guidance on comparing soil contamination with a critical concentration.
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Chemical and data (mg/kg)	STATISTICAL ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA Mean Absolute Deviation for potential outliers 3.5 = critical value of test statistic Note - MAD not applicable as 50% or more of values are the same.		
(blue denotes <= RL) (red denotes >= GAC)			
Benzo(a)pyrene	Potential	Sample	 Note - MAD not applicable as 50% or more of values are the same. Visual assessment - Q-g & histogram plots
	Outlier?	-	Q-q plot
0.01	n/a n/a	HTP07 @ 0.50 HTP09 @ 0.20	
	n/a	HTP10 @ 0.50	
0.01	}		
0.01	n/a	HTP13 @ 0.20	
			Sample (vi)
			du -3 -2 -1 0 1 2 3
			-2
			-3 Theoretical (zi)
			·····
			Benzo(a)pyrene
			3.5
			3 <u>\$</u> 2.5
			2.5 2 1.5 0.5 0.5 0.01 0.01 0.01 0.01 0.01 0.
			Concentration
			Data Normal distribution
			Basic data Risk parameter Human health - residential with plant uptake (2.5%SOM)
			0.94 = GAC (critical conc.) (mg/kg) 4 = no. samples 0 = no. samples > or = GAC
			0.01 = max. value 0.01 = max. value 0.01 = laboratory reporting limit (RL)
			0.01 = mean 4 = no. samples at RL
			Statistical tests One-sample t-test One-sided Chebychev Theorem
			$\begin{array}{c c} -3.72E+12 = t_0 & -3.7E+12 = k_0 \\ 2.353 = t_{(n-1,0.95)} & 4.36 = k_{0.05} \end{array}$
			0.01 = 95% UCL (US ₉₅) $0.01 = 95%$ UCL (US ₉₅)
			Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H_1) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC US ₉₅ = 0.01 GAC = 0.94 (US95 = 0.011 × GAC)
			Site reference POTENTIALLY SUITABLE FOR USE Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.





Chemical and data (mg/kg)	STATISTICAL ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA			
(blue denotes <= RL)		Mean Absolute Deviation for potential outliers 3.5 = critical value of test statistic Note - MAD not applicable as 50% or more of values are the same		
(red denotes >= GAC) Benzo(ghi)perylene	Potential	Sample	est statistic Note - MAD not applicable as 50% or more of values are the same. Visual assessment - Q-g & histogram plots	
0.01	Outlier? n/a	HTP07 @ 0.50	Q-q plot	
0.01	n/a	HTP09 @ 0.20		
0.01	n/a	HTP10 @ 0.50		
0.01	n/a	HTP13 @ 0.20	3	
0.01	11/a	1111 13 @ 0.20		
			2	
			- <u>-</u>	
			Sample (yi)	
			-1 -1 -1 -1 -1	
			-2	
			_	
			Theoretical (zi)	
			Benzo(ghi)perylene	
			3.5	
			3	
			- <u>\$</u> 2.5	
			2.5 2 1.5 0 0 0.5 0 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01	
			ح ح	
			- Z 0.01 0.01 0.01 0.01 0.01 0.01	
			Concentration	
			Data Normal distribution	
			Basic data Risk parameter	
			Human health - residential with plant uptake (2.5%SOM)	
			46 = GAC (critical conc.) (mg/kg)	
			4 = no. samples $0 = no. samples > or = GAC$	
			0.01 = min. value	
			0.01 = max. value 0.01 = laboratory reporting limit (RL)	
			0.01 = mean 4 = no. samples at RL	
			5E-13 = standard deviation RL is limit of detection of the method used	
			Statistical tests	
			One-sample t-test One-sided Chebychev Theorem	
			$-1.84E+14 = t_0$ $-1.8E+14 = k_0$	
			$2.353 = t_{(n-1,0.95)}$ $4.36 = k_{0.05}$	
			0.01 = 95% UCL (US ₉₅) $0.01 = 95%$ UCL (US ₉₅)	
			Results of significance test at 95% confidence level	
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC	
			Alternative hypotheses (H_1) = level of contamination is lower than the GAC	
		-	Data set treated as non-normally distributed	
			Therefore:	
			Use Chebychev Theorem - Ho rejected, true mean <=GAC	
			$US_{95} = 0.01$ GAC = 46 (US95 = 0 x GAC)	
			Site reference POTENTIALLY SUITABLE FOR USE	
			Data set: CMF & DF	
			Client: Bovis Barratt and Taylor Wimpy	
			Site: Land at Bankside, Banbury	
			Job no: C12702	
			Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.	
	1	1	Autorence. Or Anne & Oren, may 2000. Outdance on companing son contamination with a childer concentration.	



Chemical and data (mg/kg)		STATISTICAL	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL) (red denotes >= GAC)	3.5	lute Deviation for p = critical value of te	
Benzo(k)fluoranthene	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.01	n/a	HTP07 @ 0.50	<u>Q-q plot</u>
0.01	n/a	HTP09 @ 0.20	
0.01	n/a	HTP10 @ 0.50	3
0.01	n/a	HTP13 @ 0.20	
			Sample (y)
			$ \mathbf{r}$ \mathbf{r} r
			-2
			Theoretical (zi)
			Benzo(k)fluoranthene
			<u>Denzo(k)/nuorantnene</u>
			3.5
			3
			- £ 2.5
			ste 2.5 1.5 0 0.5 0 0.01 0.01 0.01 0.01
			- ō
			Concentration
			~~~~
			Data      Normal distribution
			****
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			9.6 = GAC (critical conc.) (mg/kg)
			4 = no.  samples $0 = no.  samples > or = GAC$
			0.01 = min. value
			0.01 = max. value 0.01 = laboratory reporting limit (RL) 0.01 = mean 4 = no. samples at RL
			0.01 = mean 4 = no. samples at RL 5E-13 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-3.84E+13 = t_0$ -3.8E+13 = $k_0$
			$\begin{array}{c} 2.353 = t_{(n-1,0.95)} \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) \\ 0.01 = 0.55(-1.01) $
			0.01 = 95% UCL (US ₉₅ ) $0.01 = 95%$ UCL (US ₉₅ )
			Results of significance test at 95% confidence level
			Null hypothesis ( $H_0$ ) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
		1	Use Chebychev Theorem - Ho rejected, true mean <=GAC US ₉₅ = 0.01 GAC = 9.6 (US95 = 0.001 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			Job no: C12702
	1		Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.

Chemical and data (mg/kg)	STATISTICAL ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA		
(blue denotes <= RL) (red denotes >= GAC)		olute Deviation for po = critical value of te	
Chrysene	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.013	Outlier	HTP07 @ 0.50	Q-q plot
0.01		HTP09 @ 0.20	
0.01		HTP10 @ 0.50	
0.012		HTP13 @ 0.20	
0.012		1111 10 @ 0.20	
			2
	-		
			Beneficial State S
			-2
			_
			Theoretical (zi)
			Chrysene
			_
			2.5
			·····
			2
			1.5 1.5 0 0 0 0 0.005 0.01 0.015 0.02
			- ⁷ 0 0.5
	-		<b>ž</b> 0 0.005 0.01 0.015 0.02
			Concentration
			Data      Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			8 = GAC (critical conc.) (mg/kg)
			4 = no.  samples $0 = no.  samples > or = GAC$
			0.01 = min. value
			0.013 = max. value 0.01 = laboratory reporting limit (RL)
			0.01125 = mean 2 = no. samples at RL
			0.0015 = standard deviation RL is limit of detection of the method used
			Statistical tasts
			Statistical tests One-sample t-test One-sided Chebychev Theorem
			$-10651.67 = t_0$ $-10651.7 = k_0$
			$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			0.013015 = 95% UCL (US ₉₅ ) $0.01452 = 95%$ UCL (US ₉₅ )
			Results of significance test at 95% confidence level
			Null hypothesis $(H_0)$ = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			US ₉₅ = 0.01452 GAC = 8 (US95 = 0.002 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			Job no: C12702
	1	1	Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.

Chemical and data (mg/kg) (blue denotes <= RL)	Mean Abso	STATISTICAL	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(red denotes >= GAC)	3.5	= critical value of te	
Dibenz(a,h)anthracene	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.01 0.01	n/a n/a	HTP07 @ 0.50 HTP09 @ 0.20	<u>Q-q plot</u>
0.01	n/a	HTP10 @ 0.50	
0.01	n/a	HTP13 @ 0.20	3
			2
			Sample (j)
			-2
			Theoretical (zi)
			Dibenz(a,h)anthracene
			3.5
			3
			ste 2.5 1.5 0 0.5 0 0.1 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01
			Z 0.01 0.01 0.01 0.01 0.01
			Concentration
			Concentration
			Data      Normal distribution
			Basic data Risk parameter
		1	Human health - residential with plant uptake (2.5%SOM) 0.86 = GAC (critical conc.) (mg/kg)
			4 = no. samples 0 = no. samples > or = GAC
			0.01 = min. value
			0.01 = max. value 0.01 = laboratory reporting limit (RL) 0.01 = mean 4 = no. samples at RL
			5E-13 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-3.4E+12 = t_0$ $-3.4E+12 = k_0$
			$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			$0.01 = 95\% \text{ UCL } (\text{US}_{95})$ $0.01 = 95\% \text{ UCL } (\text{US}_{95})$
			Results of significance test at 95% confidence level
			Null hypothesis ( $H_0$ ) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			US ₉₅ = 0.01 GAC = 0.86 (US95 = 0.012 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.



Chemical and data (mg/kg) (blue denotes <= RL)	Mean Abs	STATISTICAL olute Deviation for p	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(red denotes >= GAC)		5 = critical value of te	
Fluoranthene	Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.047		HTP07 @ 0.50	<u>Q-q plot</u>
0.01		HTP09 @ 0.20	
0.031		HTP10 @ 0.50	
0.088		HTP13 @ 0.20	
			2
			Sample (yi)
			-2
	i		
			Theoretical (zi)
			3.5
			<b>£</b> 2.5
			2.5 2 1.5 0.6 0.05 0.1 0.15 0.2
			<b>ž</b> -0.15 -0.1 -0.05 0 0.05 0.1 0.15 0.2
			Concentration
			Data      Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			460 = GAC (critical conc.) (mg/kg)
			4 = no.  samples $0 = no.  samples > or = GAC$
			0.01 = min. value 0.088 = max. value 0.01 = laboratory reporting limit (RL)
			0.044 = mean $1 = no. samples at RL$
			0.033015 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test         One-sided Chebychev Theorem           -27863.33 = t ₀ -27863.3 = k ₀
			$0.082842 = 95\% \text{ UCL (US}_{95}) \qquad 0.115973 = 95\% \text{ UCL (US}_{95})$
			Results of significance test at 95% confidence level
			Null hypothesis $(H_0)$ = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean $\leq$ GAC USec = 0.115973 GAC = 460 (US95 = 0.2 GAC)
			US ₉₅ = 0.115973         GAC = 460         (US95 = 0 x GAC)           Site reference         POTENTIALLY SUITABLE FOR USE
			Site reference POTENTIALLY SUITABLE FOR USE
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.



Chemical and data (mg/kg)	STATISTICAL ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA		
(blue denotes <= RL)		olute Deviation for po	
(red denotes >= GAC)	Potential	5 = critical value of te Sample	st statistic Note - MAD not applicable as 50% or more of values are the same. Visual assessment - Q-q & histogram plots
0.01	Outlier? n/a	HTP07 @ 0.50	Q-q plot
0.01	n/a	HTP09 @ 0.20	
0.01	n/a	HTP10 @ 0.50	
0.01	n/a	HTP13 @ 0.20	3
			2
			Sample (j) Sample (j) Sample (j)
			-2
			Theoretical (zi)
			Fluorene
			3.5
			- 22 2.5
			ق
			steme     2       1.5     1.5       0     0.5       0     0.01       0.01     0.01
			<b>ž</b> 0.01 0.01 0.01 0.01 0.01 0.01
			Concentration
			Data Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			380 = GAC (critical conc.) (mg/kg)         4 = no. samples       0 = no. samples > or = GAC
			- $0.01 = min. value$
			0.01 = max. value 0.01 = laboratory reporting limit (RL)
			0.01 = mean 4 = no. samples at RL
			5E-13 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sided Chebychev Theorem
			$-1.52E+15 = t_0$ $-1.5E+15 = k_0$
			$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			0.01 = 95% UCL (US ₉₅ ) $0.01 = 95%$ UCL (US ₉₅ )
			Results of significance test at 95% confidence level
			Null hypothesis ( $H_0$ ) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses ( $H_1$ ) = level of contamination is lower than the GAC
			Data set treated as <b>non-normally distributed</b>
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			US ₉₅ = 0.01 GAC = 380 (US95 = 0 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy Site: Land at Bankside, Banbury
			Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008.Guidance on comparing soil contamination with a critical concentration.

Chemical and data (mg/kg)	STATISTICAL ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA Mean Absolute Deviation for potential outliers 3.5 = critical value of test statistic Note - MAD not applicable as 50% or more of values are the same.		
(blue denotes <= RL) (red denotes >= GAC)			
Indeno(1,2,3,cd)pyrene	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.01	n/a	HTP07 @ 0.50	<u>Q-q plot</u>
0.01	n/a	HTP09 @ 0.20	
0.01	n/a	HTP10 @ 0.50	_
0.01	n/a	HTP13 @ 0.20	3
			2
			Sample (Xi)
			Theoretical (zi)
			Indeno(1,2,3,cd)pyrene
			3.5
			3
			<b>£</b> 2.5
			2.5 2.5 1.5 0.5 0.01 0.01 0.01 0.01 0.01 0.01 0.01 0.01
			<b>2</b> 0.01 0.01 0.01 0.01 0.01 0.01
			• • • • • • • • • • • • • • • • • • •
			Concentration
			DataNormal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			3.9 = GAC (critical conc.) (mg/kg)
		+	4 = no.  samples $0 = no.  samples > or = GAC$
			4 = 10. samples $0 = 10.$ samples $> 0 = 0.01$
			0.01 = max. value $0.01 = laboratory reporting limit (RL)$
			0.01 = max. value $0.01 = nabolatory reporting mint (RL)0.01 = mean 4 = no. samples at RL$
			5E-13 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-1.56E+13 = t_0$ $-1.6E+13 = k_0$
			$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			$0.01 = 95\% \text{ UCL } (\text{US}_{95})$ $0.01 = 95\% \text{ UCL } (\text{US}_{95})$
			Results of significance test at 95% confidence level
			Null hypothesis $(H_0)$ = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			US ₉₅ = 0.01 GAC = 3.9 (US95 = 0.003 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			Job no: C12702
	1		Reference: CL:AIRE & CIEH. May 2008.Guidance on comparing soil contamination with a critical concentration.
		1	

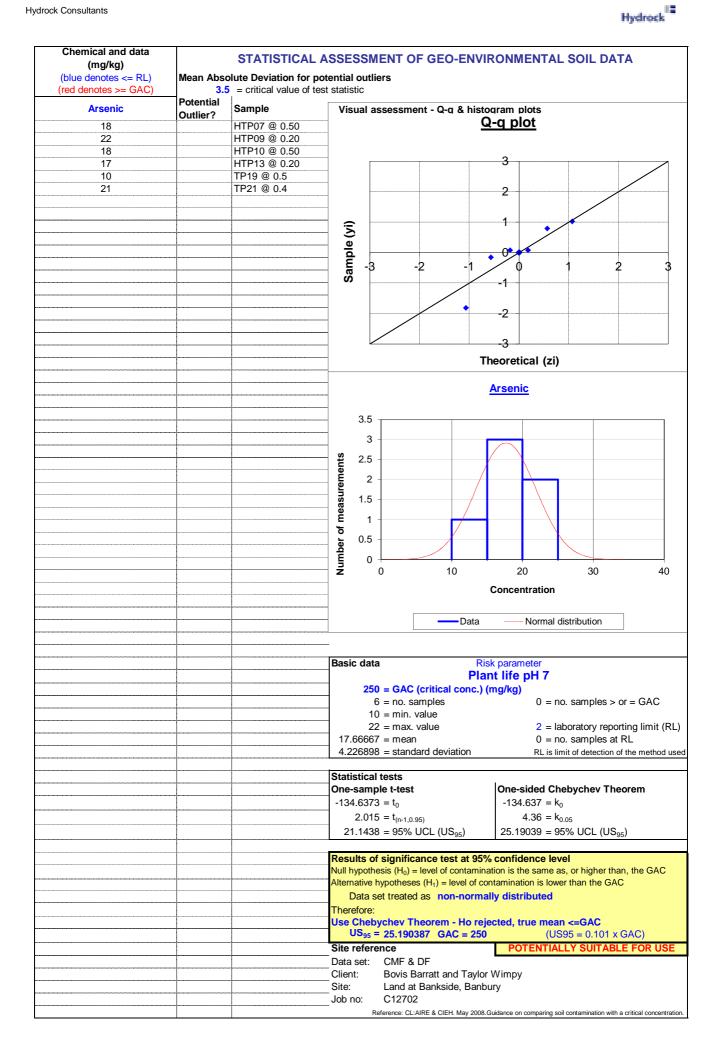


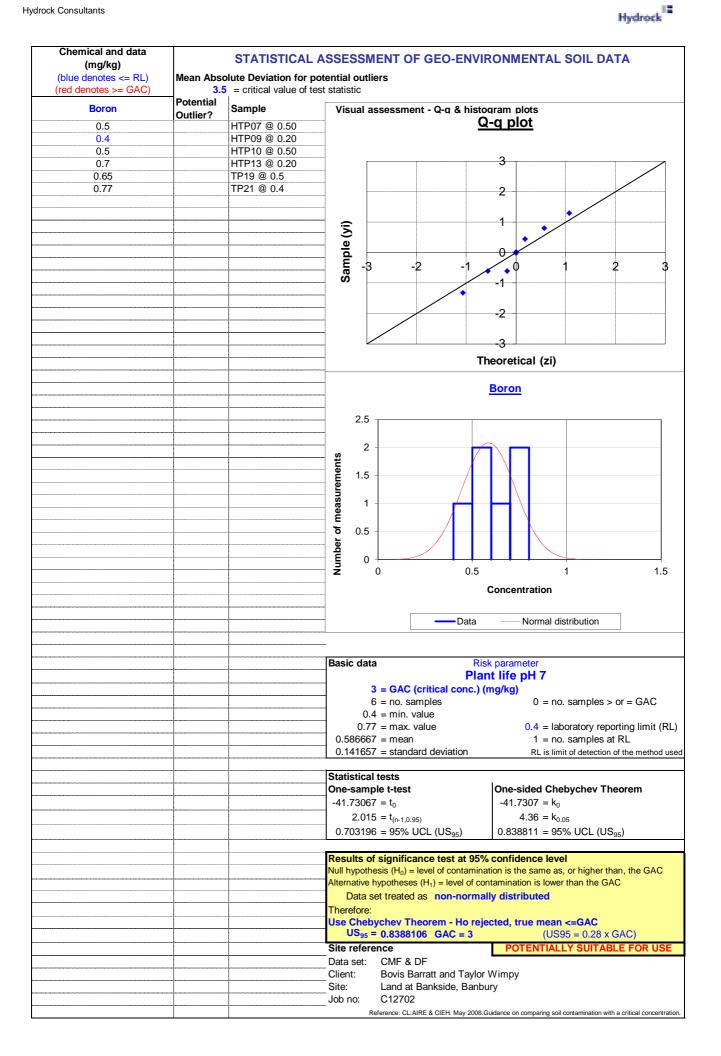
Chemical and data (mg/kg)	STATISTICAL ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA Mean Absolute Deviation for potential outliers		
(blue denotes <= RL)			
(red denotes >= GAC) Naphthalene	Potential	5 = critical value of te Sample	<ul> <li>Statistic Note - MAD not applicable as 50% or more of values are the same.</li> <li>Visual assessment - Q-g &amp; histogram plots</li> </ul>
•	Outlier?	•	
0.01	n/a	HTP07 @ 0.50 HTP09 @ 0.20	
0.01	n/a	HTP10 @ 0.50	
	n/a		
0.01	n/a	HTP13 @ 0.20	
			2
			Beneficial Sector Secto
			S 1
			-1
			-2
			Theoretical (zi)
			Naphthalene
			3.5
			3
			- <b>£</b> 2.5
			0.5
			2.5 2 1.5 0 0.5 0 0.01 0.01 0.01 0.01 0.01 0.
			- <b>N</b> 0.01 0.01 0.01 0.01 0.01 0.01
			Concentration
			••••
			Data      Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			3.7 = GAC (critical conc.) (mg/kg)
			4 = no.  samples $0 = no.  samples  > or = GAC$
			0.01 = min. value
			0.01 = max. value 0.01 = laboratory reporting limit (RL)
			$0.01 = mean \qquad 4 = no. samples at RL$
			5E-13 = standard deviation RL is limit of detection of the method used
			Statistical tools
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-1.48E+13 = t_0$ $-1.5E+13 = k_0$
			$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			0.01 = 95% UCL (US ₉₅ ) $0.01 = 95%$ UCL (US ₉₅ )
			Results of significance test at 95% confidence level
			Null hypothesis $(H_0)$ = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			US ₉₅ = 0.01 GAC = 3.7 (US95 = 0.003 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.
•			

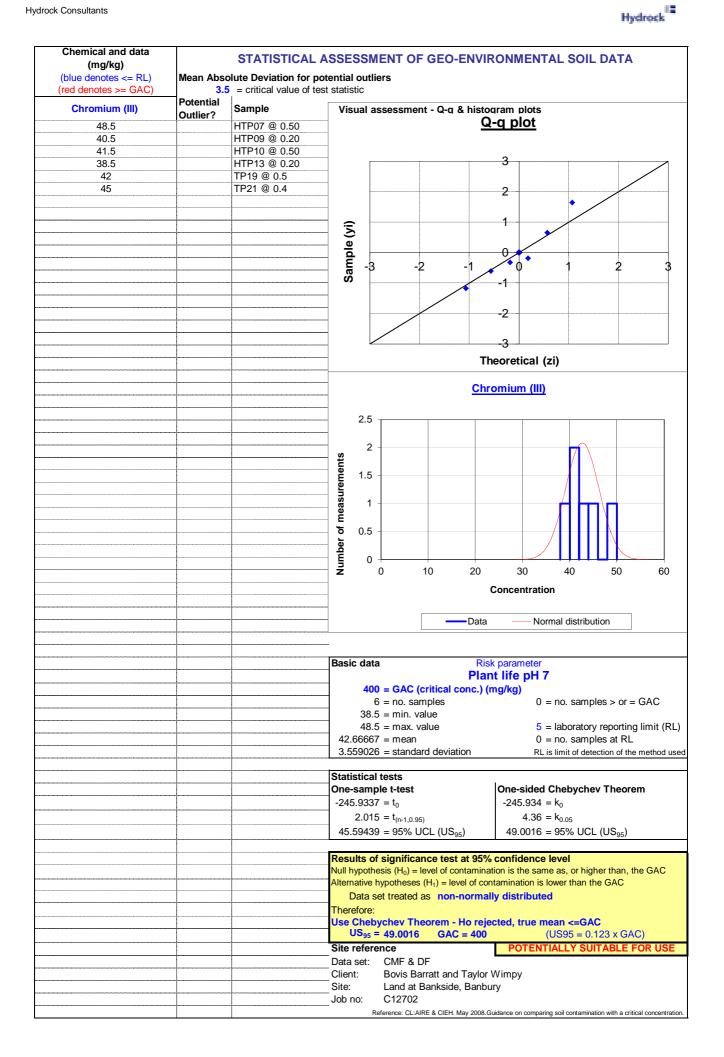
Chemical and data (mg/kg)		STATISTICAL A	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL) (red denotes >= GAC)		<b>blute Deviation for pot</b>	
Phenanthrene	Potential	Sample	Visual assessment - Q-q & histogram plots
0.018	Outlier?	HTP07 @ 0.50	Q-q plot
0.018		HTP07 @ 0.50	
0.01		HTP10 @ 0.50	-
0.014		HTP13 @ 0.20	3
			2
			-2
			Theoretical (zi)
			 Phenanthrene
			3.5
			3
			- <b>st</b>
			1.5
			2.5 2 1.5 0.5 -0.01 0 0.01 0.02 0.03
			<b>ž</b> -0.01 0 0.01 0.02 0.03
			Concentration
			Data      Normal distribution
			M
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			200 = GAC (critical conc.) (mg/kg)
			4 = no. samples 0 = no. samples > or = GAC
			0.01 = min. value 0.018 = max. value 0.01 = laboratory reporting limit (RL)
			0.013 = mean $2 = no. samples at RL$
			0.00383 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sided Chebychev Theorem
			$-104439.8 = t_0$ $-104440 = k_0$
			$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			$0.017506 = 95\% \text{ UCL } (\text{US}_{95})$ $0.021349 = 95\% \text{ UCL } (\text{US}_{95})$
			Depute of similiance test of 050/ acrifichence level
			<b>Results of significance test at 95% confidence level</b> Null hypothesis ( $H_0$ ) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses ( $H_1$ ) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC US ₉₅ = 0.0213488 GAC = 200 (US95 = 0 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury Job no: C12702
			JOD NO: C12/U2 Reference: CL:AIRE & CIEH. May 2008.Guidance on comparing soil contamination with a critical concentration.
	1	1	reference. Service a oren in may 2000, Suidance on comparing Son Conditionation with a Chical Concentration.



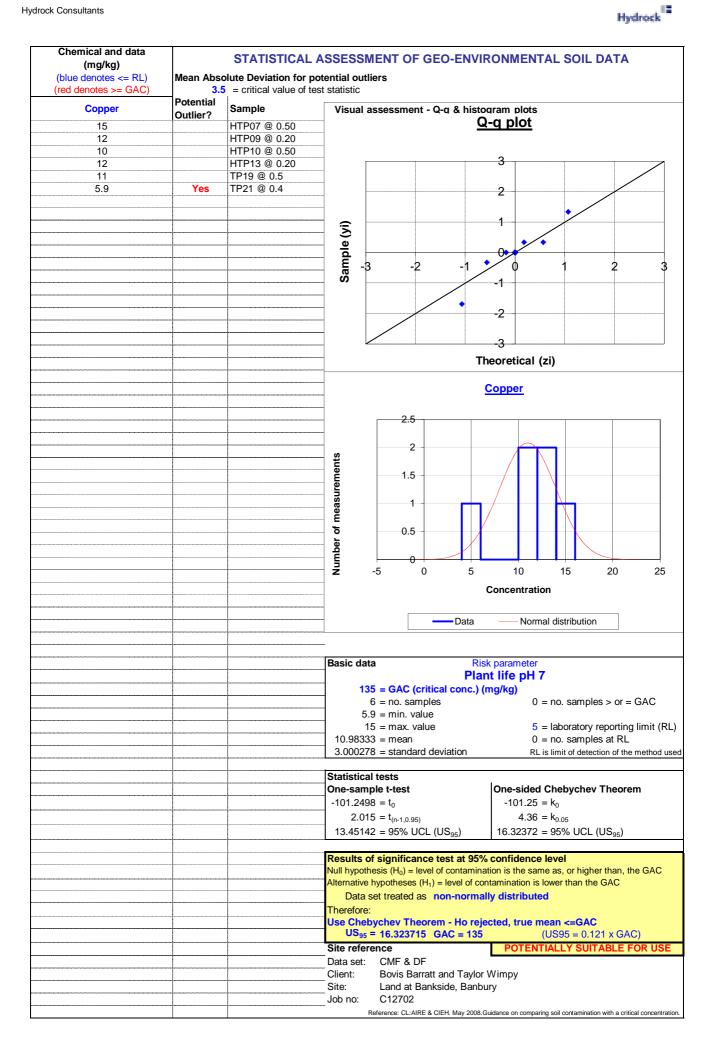
Chemical and data (mg/kg) (blue denotes <= RL)		olute Deviation for po	
(red denotes >= GAC)	3.5 Potential	= critical value of te	
Pyrene	Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.035		HTP07 @ 0.50	<u>Q-q plot</u>
0.01		HTP09 @ 0.20	
0.017		HTP10 @ 0.50	-
0.061		HTP13 @ 0.20	3
			2
			$-\overline{5}$
			Sample (j)
			-2
			<b>.</b>
			Theoretical (zi)
			_
			Pyrene
			2.5
			1.5
			- <del>5</del> 0.5
			1.5 0.5 0.0 0.05 0.1 0.15
			<b>- - - 0.1</b> -0.05 0 0.05 0.1 0.15
			Concentration
			Concentration
			Data Normal distribution
			····
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			1000 = GAC (critical conc.) (mg/kg)
			4 = no. samples 0 = no. samples > or = GAC
			0.01 = min. value 0.061 = max. value 0.01 = laboratory reporting limit (RL)
			0.03075 = mean 0.03075 = nean 1 = no. samples at RL
			0.02275 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-87907.62 = t_0$ $-87907.6 = k_0$
			$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			0.057516 = 95% UCL (US ₉₅ ) $0.080346 = 95%$ UCL (US ₉₅ )
			Results of significance test at 95% confidence level
			Null hypothesis $(H_0)$ = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			$US_{95} = 0.080346$ GAC = 1000 (US95 = 0 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF
			Client: Bovis Barratt and Taylor Wimpy
			_Site: Land at Bankside, Banbury Job no: C12702
			JOD NO. C12702 Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.
	1	1	Nererence, OLIANNE & OLET IN WAY 2000. Outdative on companing soil contamination with a critical concentration.

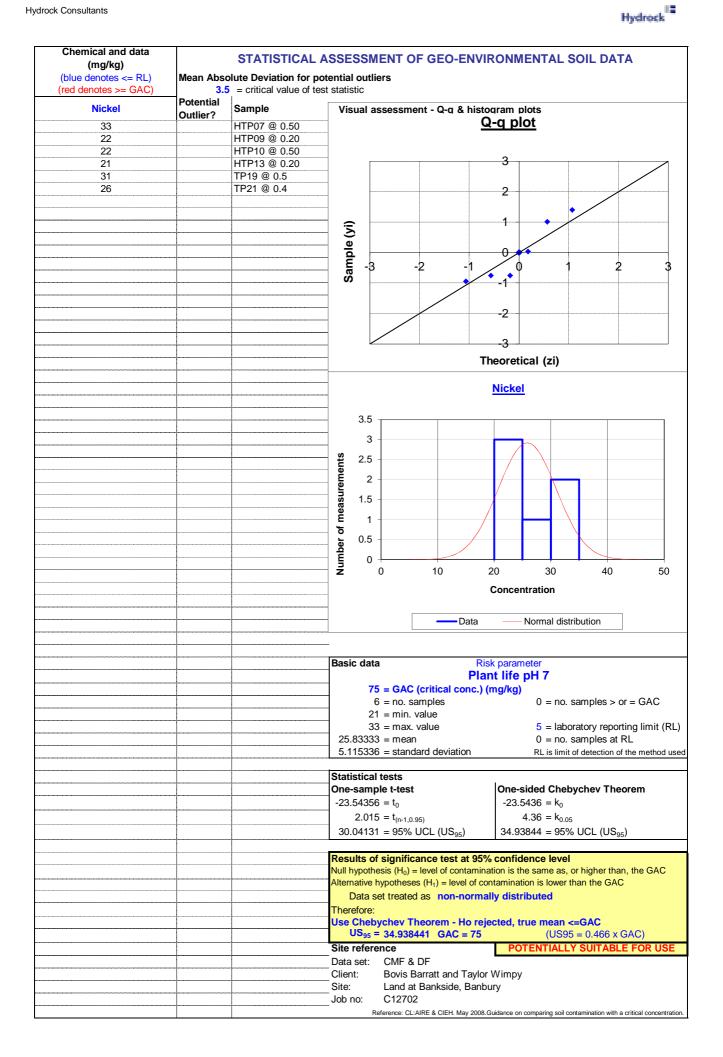


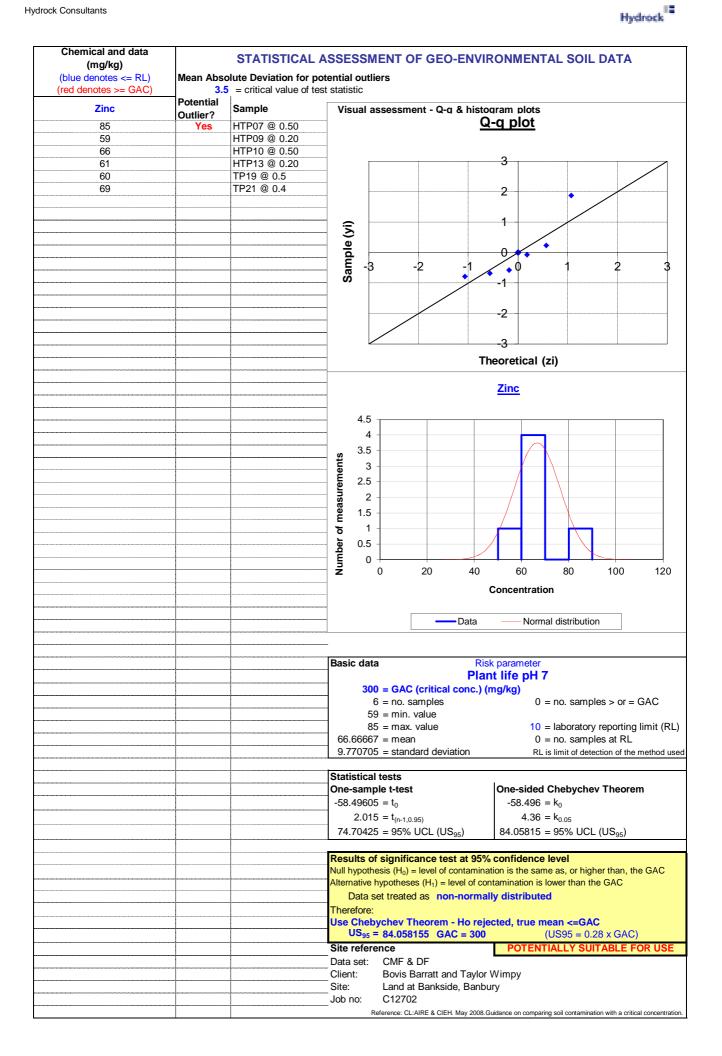




Chemical and data (mg/kg)		STATISTICAL	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL) (red denotes >= GAC)		<b>blute Deviation for p</b>	
Chromium (VI)	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.5	n/a	HTP07 @ 0.50	<u>Q-q plot</u>
0.5	n/a	HTP09 @ 0.20	
0.5	n/a n/a	HTP10 @ 0.50 HTP13 @ 0.20	3
0.5	II/a	1117 13 @ 0.20	
			2
			Sample (y)
			-1-1
			-2
			Theoretical (zi)
			Chromium (VI)
			3.5
			3
			<b>£</b> 2.5
			- ⁷ 0 - 0.5
			<b>Z</b> 0.5 0.5 0.5 0.5 0.5 0.5
			Concentration
			Data — Normal distribution
			Basic data Risk parameter Plant life pH 7
			25 = GAC (critical conc.) (mg/kg)
			4 = no. samples 0 = no. samples > or = GAC
			0.5 = min. value
			0.5 = max. value 0.5 = laboratory reporting limit (RL) 0.5 = mean 4 = no. samples at RL
			5E-12 = standard deviation RL is limit of detection of the method used
			Chasing in the sector
			Statistical tests One-sample t-testOne-sided Chebychev Theorem
			$-9.8E+12 = t_0$ $-9.8E+12 = k_0$
			$2.353 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			0.5 = 95% UCL (US ₉₅ ) $0.5 = 95%$ UCL (US ₉₅ )
			Results of significance test at 95% confidence level
			Null hypothesis ( $H_0$ ) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			US ₉₅ = 0.5 GAC = 25 (US95 = 0.02 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: CMF & DF Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.









1								Soil Type	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT
	All values	in mg/kg unle	ess otherwi	se stated				Location & Depth	HTP15	HTP17	HTP20	HTP22	HTP24	HTP25	HTP26	HTP29	HTP33	HTP39	HTP40
Chemical of Potential Concern	Lab. RL	No. Samples	Min. Value	Max. Value	No. Samples > or = GAC	GAC	US ₉₅	Result of Significance Test	0.25	0.15	0.15	0.20	0.20	0.15	0.10	0.15	0.25	0.20	0.20
Arsenic	2	40	17	230	37	32	110.5811	FURTHER ASSESSMENT REQUIRED	110	66	68	86	79	140	120	58	76	52	58
Beryllium	1	18	0.5	5.6	0	51	4.442717	POTENTIALLY SUITABLE FOR USE	4.1	1.6	2.3	2.5	2	4.2	3.7	1.6	2.1	1.4	1.6
Boron	0.4	20	0.5	2.2	0	290	1.440611	POTENTIALLY SUITABLE FOR USE	0.8	0.9	1.4	0.7	0.9	0.5	1.1	1.3	1.2	0.7	0.9
Cadmium	0.1	23	0.1	1.8	0	11	0.693381	POTENTIALLY SUITABLE FOR USE	0.1	0.1	0.28	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Chromium (III)	5	23	46.6	589.5	0	630	329.0309	POTENTIALLY SUITABLE FOR USE	269.5	77.5	87.5	109.5	109.5	269.5	239.5	77.5	99.5	77.5	81.5
Chromium (VI)	0.5	20	0.2	1.2	0	4.3	0.683627	POTENTIALLY SUITABLE FOR USE	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
Copper	5	23	6	49	0	2300	38.32559	POTENTIALLY SUITABLE FOR USE	21	23	31	31	31	41	46	28	40	23	26
Lead	5	23	19	120	0	450	82.2043	POTENTIALLY SUITABLE FOR USE	39	46	62	55	51	70	120	48	71	44	49
Mercury, inorganic	0.1	23	0.1	0.6	0	170	0.349307	POTENTIALLY SUITABLE FOR USE	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1	0.1
Nickel	5	40	29	210	8	130	110.2809	POTENTIALLY SUITABLE FOR USE	110	62	99	100	92	150	120	69	79	52	60
Selenium	0.2	23	0.2	3	0	350	1.454144	POTENTIALLY SUITABLE FOR USE	0.2	0.2	0.2	0.2	0.2	0.22	0.2	0.2	0.2	0.2	0.2
Vanadium	5	16	100	740	16	74	549.9117	FURTHER ASSESSMENT REQUIRED	410	120	150	150	160	380	320	120	150	100	120
Zinc	10	23	74	970	0	3700	393.701	POTENTIALLY SUITABLE FOR USE	190	110	180	140	160	300	260	110	150	100	120
Cyanide (free)	0.5	20	0.5	5	0	750	3.200455	POTENTIALLY SUITABLE FOR USE	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
Phenol (total)	0.3	20	0.01	1.1	0	290	0.539339	POTENTIALLY SUITABLE FOR USE	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3	0.3
Acenaphthene	0.01	20	0.01	0.5	0	480	0.312093	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01
Acenaphthylene	0.01	20	0.01	0.5	0	400	0.309073	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.073	0.026
Anthracene	0.01	20	0.01	0.78	0	4900	0.363842	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.11	0.01	0.01	0.01	0.01	0.012	0.01	0.14	0.064
Benz(a)anthracene	0.01	20	0.01	0.95	0	4.7	0.507781	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.36	0.01	0.01	0.01	0.047	0.099	0.023	0.95	0.48
Benzo(a)pyrene	0.01	20	0.01	0.95	1	0.94	0.483634	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.38	0.01	0.01	0.01	0.01	0.07	0.018	0.69	0.33
Benzo(b)fluoranthene	0.01	20	0.01	1.2	0	6.5	0.575744	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.5	0.01	0.01	0.01	0.046	0.12	0.021	1.2	0.62
Benzo(ghi)perylene	0.01	20	0.01	0.5	0	46	0.320275	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.12	0.01	0.01	0.01	0.01	0.013	0.01	0.21	0.076
Benzo(k)fluoranthene	0.01	20	0.01	0.5	0	9.6	0.333785	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.13	0.01	0.01	0.01	0.01	0.011	0.01	0.35	0.099
Chrysene	0.01	20	0.01	0.78	0	8	0.454279	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.3	0.01	0.01	0.01	0.029	0.08	0.024	0.78	0.32
Dibenz(a,h)anthracene	0.01	20	0.01	0.5	0	0.86	0.30405	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01
Fluoranthene	0.01	20	0.01	2.3	0	460	1.011848	POTENTIALLY SUITABLE FOR USE	0.096	0.019	0.81	0.043	0.01	0.028	0.13	0.26	0.076	1.7	0.84
Fluorene	0.01	20	0.01	0.5	0	380	0.311894	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01
Indeno(1,2,3,cd)pyrene	0.01	20	0.01	0.5	0	3.9	0.318916	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.083	0.01	0.01	0.01	0.01	0.01	0.01	0.22	0.069
Naphthalene	0.01	20	0.01	2	0	3.7	1.049686	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01	0.01
Phenanthrene	0.01	20	0.01	2.1	0	200	0.725232	POTENTIALLY SUITABLE FOR USE	0.039	0.01	0.39	0.01	0.01	0.01	0.043	0.059	0.029	0.28	0.17
Pyrene	0.01	20	0.01	2.3	0	1000	0.928259	POTENTIALLY SUITABLE FOR USE	0.037	0.028	0.64	0.03	0.01	0.024	0.1	0.21	0.056	1.4	0.65
	Mean																		
FOC (dimensionless)	0.020								0.0099	0.017	0.023	0.021	0.057	0.012	0.013	0.023	0.017	0.011	0.013
SOM (calculated)	3.40%								1.71%	2.93%	3.97%	3.62%	9.83%	2.07%	2.24%	3.97%	2.93%	1.90%	2.24%
pH (su)	7.7								7.1	7.5	7.7	8.3	7.7	7.7	7.9	7.9	7.6	7.6	7.3

Risk parameter: Human health - residential with plant uptake (2.5%SOM)

Data set: MRB & WMF Client: Bovis Barratt and Taylor Wimpy

Site: Land at Bankside, Banbury

Job no: C12702

Legend: Values in blue are at or below the laboratory reporting limit (where a single value is indicated) and are considered as being at the detection limit for the purposes of statistical analysis, as a conservative es Values in red are equal to, or greater than, the generic assessment criterion (GAC).

MG denotes Made Ground

NAT denotes natural ground

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								Soil Type	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT
	All values	in mg/kg unle	ess otherwi	se stated				Location & Depth	HTP41	HTP43	HTP44	HTP45	HTP46	TP5	TP8	TP8	TP9	HS1 TS ⁴
Chemical of Potential Concern	Lab. RL	No. Samples	Min. Value	Max. Value	No. Samples > or = GAC	GAC	US ₉₅	Result of Significance Test	0.1	0.2	0.7	0.4	0.7	0.1	0.4	2.8	0.6	0.21
Arsenic	2	40	17	230	37	32	110.5811	FURTHER ASSESSMENT REQUIRED	190	130	230	120	170	19	58	17	27	120
Beryllium	1	18	0.5	5.6	0	51	4.442717	POTENTIALLY SUITABLE FOR USE	5.4	3.6	5.6	3.3	4.8	0.5	0.5			
Boron	0.4	20	0.5	2.2	0	290	1.440611	POTENTIALLY SUITABLE FOR USE	1	1	0.7	1	1.2	1.8	1	2.2	0.7	
Cadmium	0.1	23	0.1	1.8	0	11	0.693381	POTENTIALLY SUITABLE FOR USE	0.36	0.37	0.27	0.4	0.23	0.5	0.5	0.5	0.5	
Chromium (III)	5	23	46.6	589.5	0	630	329.0309	POTENTIALLY SUITABLE FOR USE	479.5	209.5	539.5	189.5	589.5	55.57	109.8	53.8	46.6	
Chromium (VI)	0.5	20	0.2	1.2	0	4.3	0.683627	POTENTIALLY SUITABLE FOR USE	0.5	0.5	0.5	0.5	0.5	0.43	0.2	1.2	0.4	
Copper	5	23	6	49	0	2300	38.32559	POTENTIALLY SUITABLE FOR USE	40	49	35	43	25	11	18	13	12	
Lead	5	23	19	120	0	450	82.2043	POTENTIALLY SUITABLE FOR USE	81	90	84	81	33	32	37	25	19	
Mercury, inorganic	0.1	23	0.1	0.6	0	170	0.349307	POTENTIALLY SUITABLE FOR USE	0.19	0.19	0.13	0.17	0.1	0.2	0.27	0.2	0.2	
Nickel	5	40	29	210	8	130	110.2809	POTENTIALLY SUITABLE FOR USE	170	160	210	130	180	29	57	32	43	100
Selenium	0.2	23	0.2	3	0	350	1.454144	POTENTIALLY SUITABLE FOR USE	0.2	0.2	0.2	0.2	0.2	0.3	0.3	0.3	0.3	
Vanadium	5	16	100	740	16	74	549.9117	FURTHER ASSESSMENT REQUIRED	670	330	710	290	740					
Zinc	10	23	74	970	0	3700	393.701	POTENTIALLY SUITABLE FOR USE	340	290	380	220	970	82	130	74	84	
Cvanide (free)	0.5	20	0.5	5	0	750	3.200455	POTENTIALLY SUITABLE FOR USE	0.5	0.5	0.5	0.5	0.5	5	5	5	5	
Phenol (total)	0.3	20	0.01	1.1	0	290	0.539339	POTENTIALLY SUITABLE FOR USE	0.01	0.1	0.044	0.054	0.091	0.5	0.5	1.1	0.5	
Acenaphthene	0.01	20	0.01	0.5	0	480	0.312093	POTENTIALLY SUITABLE FOR USE	0.01	0.1	0.044	0.054	0.091	0.5	0.5	0.5	0.5	
Acenaphthylene	0.01	20	0.01	0.5	0	400	0.309073	POTENTIALLY SUITABLE FOR USE	0.01	0.063	0.03	0.028	0.01	0.5	0.5	0.5	0.5	
Anthracene	0.01	20	0.01	0.78	0	4900	0.363842	POTENTIALLY SUITABLE FOR USE	0.03	0.02	0.01	0.01	0.031	0.5	0.78	0.5	0.5	
Benz(a)anthracene	0.01	20	0.01	0.95	0	4.7	0.507781	POTENTIALLY SUITABLE FOR USE	0.11	0.04	0.01	0.01	0.01	0.5	0.76	0.5	0.5	
Benzo(a)pyrene	0.01	20	0.01	0.95	1	0.94	0.483634	POTENTIALLY SUITABLE FOR USE	0.055	0.023	0.01	0.01	0.01	0.5	0.95	0.5	0.5	
Benzo(b)fluoranthene	0.01	20	0.01	1.2	0	6.5	0.575744	POTENTIALLY SUITABLE FOR USE	0.18	0.056	0.01	0.01	0.01	0.5	0.7	0.5	0.5	
Benzo(ghi)perylene	0.01	20	0.01	0.5	0	46	0.320275	POTENTIALLY SUITABLE FOR USE	0.032	0.01	0.01	0.01	0.01	0.5	0.5	0.5	0.5	
Benzo(k)fluoranthene	0.01	20	0.01	0.5	0	9.6	0.333785	POTENTIALLY SUITABLE FOR USE	0.014	0.023	0.01	0.01	0.01	0.5	0.5	0.5	0.5	
Chrysene	0.01	20	0.01	0.78	0	8	0.454279	POTENTIALLY SUITABLE FOR USE	0.11	0.054	0.01	0.01	0.01	0.5	0.72	0.5	0.5	
Dibenz(a,h)anthracene	0.01	20	0.01	0.5	0	0.86	0.30405	POTENTIALLY SUITABLE FOR USE	0.01	0.01	0.01	0.01	0.01	0.5	0.5	0.5	0.5	
Fluoranthene	0.01	20	0.01	2.3	0	460	1.011848	POTENTIALLY SUITABLE FOR USE	0.24	0.17	0.091	0.071	0.096	0.5	2.3	0.5	0.5	
Fluorene	0.01	20	0.01	0.5	0	380	0.311894	POTENTIALLY SUITABLE FOR USE	0.01	0.08	0.038	0.053	0.11	0.5	0.5	0.5	0.5	
Indeno(1,2,3,cd)pyrene	0.01	20	0.01	0.5	0	3.9	0.318916	POTENTIALLY SUITABLE FOR USE	0.025	0.01	0.01	0.01	0.01	0.5	0.5	0.5	0.5	
Naphthalene	0.01	20	0.01	2	0	3.7	1.049686	POTENTIALLY SUITABLE FOR USE	0.8	1.5	0.95	1.2	2	0.5	1.1	0.5	0.5	
Phenanthrene	0.01	20	0.01	2.1	0	200	0.725232	POTENTIALLY SUITABLE FOR USE	0.22	0.17	0.1	0.08	0.24	0.5	2.1	0.5	0.5	
Pyrene	0.01	20	0.01	2.3	0	1000	0.928259	POTENTIALLY SUITABLE FOR USE	0.17	0.12	0.072	0.045	0.063	0.5	2.3	0.5	0.5	
	Mean			1	Ť		1		<u> </u>	<u>-</u>	1	1						
FOC (dimensionless)	0.020						-		0.023	0.018	0.012	0.022	0.0053					0.025
SOM (calculated)	3.40%			1			1		3.97%	3.10%	2.07%	3.79%	0.91%					4.31%
pH (su)	7.7						+		7.7	7.5	7.6	6.9	7.9	7.2	8.1	7.4	7.1	8.2

Risk parameter: Human health - residential with plant uptake (2.5%SOM)

Data set: MRB & WMF

stimate.

Client: Bovis Barratt and Taylor Wimpy Site: Land at Bankside, Banbury

Job no: C12702



								Soil Typ	e NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT
	All values	in mg/kg unle	ess otherwi	se stated				Location & Dept	1 HS1 TS2	HS1 SS1	HS1 SS1	HS2 TS1	HS2 TS2	HS2 SS1	HS3 TS1	HS3 TS2	HS3 TS2	HS3SS1	HS4 TS1
Chemical of Potential Concern	Lab. RL	No. Samples	Min. Value	Max. Value	No. Samples > or = GAC	GAC	US ₉₅	Result of Significance Test	0.15		No depth	0.22	0.28	0.35	0.24	0.25	No depth	0.4	0.22
Arsenic	2	40	17	230	37	32	110.5811	FURTHER ASSESSMENT REQUIRED	140	150	150	110	110	110	60	66	86	58	79
Beryllium	1	18	0.5	5.6	0	51	4.442717	POTENTIALLY SUITABLE FOR USE											
Boron	0.4	20	0.5	2.2	0	290	1.440611	POTENTIALLY SUITABLE FOR USE											[
Cadmium	0.1	23	0.1	1.8	0	11	0.693381	POTENTIALLY SUITABLE FOR USE											1
Chromium (III)	5	23	46.6	589.5	0	630	329.0309	POTENTIALLY SUITABLE FOR USE											(
Chromium (VI)	0.5	20	0.2	1.2	0	4.3	0.683627	POTENTIALLY SUITABLE FOR USE											
Copper	5	23	6	49	0	2300	38.32559	POTENTIALLY SUITABLE FOR USE											
Lead	5	23	19	120	0	450	82.2043	POTENTIALLY SUITABLE FOR USE											
Mercury, inorganic	0.1	23	0.1	0.6	0	170	0.349307	POTENTIALLY SUITABLE FOR USE											(
Nickel	5	40	29	210	8	130	110.2809	POTENTIALLY SUITABLE FOR USE	100	130	150	110	110	110	75	75	87	65	83
Selenium	0.2	23	0.2	3	0	350	1.454144	POTENTIALLY SUITABLE FOR USE											
Vanadium	5	16	100	740	16	74	549.9117	FURTHER ASSESSMENT REQUIRED											1
Zinc	10	23	74	970	0	3700	393.701	POTENTIALLY SUITABLE FOR USE											
Cyanide (free)	0.5	20	0.5	5	0	750	3.200455	POTENTIALLY SUITABLE FOR USE											
Phenol (total)	0.3	20	0.01	1.1	0	290	0.539339	POTENTIALLY SUITABLE FOR USE											
Acenaphthene	0.01	20	0.01	0.5	0	480	0.312093	POTENTIALLY SUITABLE FOR USE											[
Acenaphthylene	0.01	20	0.01	0.5	0	400	0.309073	POTENTIALLY SUITABLE FOR USE											
Anthracene	0.01	20	0.01	0.78	0	4900	0.363842	POTENTIALLY SUITABLE FOR USE											
Benz(a)anthracene	0.01	20	0.01	0.95	0	4.7	0.507781	POTENTIALLY SUITABLE FOR USE											
Benzo(a)pyrene	0.01	20	0.01	0.95	1	0.94	0.483634	POTENTIALLY SUITABLE FOR USE											
Benzo(b)fluoranthene	0.01	20	0.01	1.2	0	6.5	0.575744	POTENTIALLY SUITABLE FOR USE											
Benzo(ghi)perylene	0.01	20	0.01	0.5	0	46	0.320275	POTENTIALLY SUITABLE FOR USE											
Benzo(k)fluoranthene	0.01	20	0.01	0.5	0	9.6	0.333785	POTENTIALLY SUITABLE FOR USE											
Chrysene	0.01	20	0.01	0.78	0	8	0.454279	POTENTIALLY SUITABLE FOR USE											
Dibenz(a,h)anthracene	0.01	20	0.01	0.5	0	0.86	0.30405	POTENTIALLY SUITABLE FOR USE											
Fluoranthene	0.01	20	0.01	2.3	0	460	1.011848	POTENTIALLY SUITABLE FOR USE											
Fluorene	0.01	20	0.01	0.5	0	380	0.311894	POTENTIALLY SUITABLE FOR USE											
Indeno(1,2,3,cd)pyrene	0.01	20	0.01	0.5	0	3.9	0.318916	POTENTIALLY SUITABLE FOR USE											
Naphthalene	0.01	20	0.01	2	0	3.7	1.049686	POTENTIALLY SUITABLE FOR USE											1
Phenanthrene	0.01	20	0.01	2.1	0	200	0.725232	POTENTIALLY SUITABLE FOR USE	1										
Pyrene	0.01	20	0.01	2.3	0	1000	0.928259	POTENTIALLY SUITABLE FOR USE											1
	Mean			1															
FOC (dimensionless)	0.020		1						0.021512	0.012791		0.024419		0.026744	0.026163	0.024419		0.013372	0.025581
SOM (calculated)	3.40%								3.71%	2.21%		4.21%		4.61%	4.51%	4.21%		2.31%	4.41%
pH (su)	7.7		1				1		8.1	8.1		8.1	8.1	8.1	8	8		8	7.9

Risk parameter: Human health - residential with plant uptake (2.5%SOM)

Data set: MRB & WMF Client: Bovis Barratt and Taylor Wimpy Site: Land at Bankside, Banbury Job no: C12702

								Soil Typ	e NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT
	All values	in mg/kg unle	ess otherwi	se stated				Location & Dept	hHS4TS2	HS4 SS1	HS5 TS1	HS5 TS1	HS5TS2	WAHS1	WAHS2	WAHS4
Chemical of Potential Concern	Lab. RL	No. Samples	Min. Value	Max. Value	No. Samples > or = GAC	GAC	US ₉₅	Result of Significance Test	0.19	0.38	0.16	No depth	0.17	No depth	No depth	No depth
Arsenic	2	40	17	230	37	32	110.5811	FURTHER ASSESSMENT REQUIRED	60	70	130	110	110	150	120	69
Beryllium	1	18	0.5	5.6	0	51	4.442717	POTENTIALLY SUITABLE FOR USE								
Boron	0.4	20	0.5	2.2	0	290	1.440611	POTENTIALLY SUITABLE FOR USE		1						
Cadmium	0.1	23	0.1	1.8	0	11	0.693381	POTENTIALLY SUITABLE FOR USE						1.8	0.7	0.7
Chromium (III)	5	23	46.6	589.5	0	630	329.0309	POTENTIALLY SUITABLE FOR USE		1				270	150	87
Chromium (VI)	0.5	20	0.2	1.2	0	4.3	0.683627	POTENTIALLY SUITABLE FOR USE								
Copper	5	23	6	49	0	2300	38.32559	POTENTIALLY SUITABLE FOR USE		1				6	6	14
Lead	5	23	19	120	0	450	82.2043	POTENTIALLY SUITABLE FOR USE						46	110	49
Mercury, inorganic	0.1	23	0.1	0.6	0	170	0.349307	POTENTIALLY SUITABLE FOR USE						0.6	0.6	0.6
Nickel	5	40	29	210	8	130	110.2809	POTENTIALLY SUITABLE FOR USE	78	86	110	120	110	110	98	79
Selenium	0.2	23	0.2	3	0	350	1.454144	POTENTIALLY SUITABLE FOR USE						3	3	3
Vanadium	5	16	100	740	16	74	549.9117	FURTHER ASSESSMENT REQUIRED								
Zinc	10	23	74	970	0	3700	393.701	POTENTIALLY SUITABLE FOR USE		1				310	290	170
Cyanide (free)	0.5	20	0.5	5	0	750	3.200455	POTENTIALLY SUITABLE FOR USE								
Phenol (total)	0.3	20	0.01	1.1	0	290	0.539339	POTENTIALLY SUITABLE FOR USE								
Acenaphthene	0.01	20	0.01	0.5	0	480	0.312093	POTENTIALLY SUITABLE FOR USE								
Acenaphthylene	0.01	20	0.01	0.5	0	400	0.309073	POTENTIALLY SUITABLE FOR USE	1							
Anthracene	0.01	20	0.01	0.78	0	4900	0.363842	POTENTIALLY SUITABLE FOR USE		1						
Benz(a)anthracene	0.01	20	0.01	0.95	0	4.7	0.507781	POTENTIALLY SUITABLE FOR USE								
Benzo(a)pyrene	0.01	20	0.01	0.95	1	0.94	0.483634	POTENTIALLY SUITABLE FOR USE								
Benzo(b)fluoranthene	0.01	20	0.01	1.2	0	6.5	0.575744	POTENTIALLY SUITABLE FOR USE		1						
Benzo(ghi)perylene	0.01	20	0.01	0.5	0	46	0.320275	POTENTIALLY SUITABLE FOR USE			1					
Benzo(k)fluoranthene	0.01	20	0.01	0.5	0	9.6	0.333785	POTENTIALLY SUITABLE FOR USE								
Chrysene	0.01	20	0.01	0.78	0	8	0.454279	POTENTIALLY SUITABLE FOR USE								
Dibenz(a,h)anthracene	0.01	20	0.01	0.5	0	0.86	0.30405	POTENTIALLY SUITABLE FOR USE	1							
Fluoranthene	0.01	20	0.01	2.3	0	460	1.011848	POTENTIALLY SUITABLE FOR USE								
Fluorene	0.01	20	0.01	0.5	0	380	0.311894	POTENTIALLY SUITABLE FOR USE	1							
Indeno(1,2,3,cd)pyrene	0.01	20	0.01	0.5	0	3.9	0.318916	POTENTIALLY SUITABLE FOR USE								
Naphthalene	0.01	20	0.01	2	0	3.7	1.049686	POTENTIALLY SUITABLE FOR USE								
Phenanthrene	0.01	20	0.01	2.1	0	200	0.725232	POTENTIALLY SUITABLE FOR USE								
Pyrene	0.01	20	0.01	2.3	0	1000	0.928259	POTENTIALLY SUITABLE FOR USE								
	Mean								1	1						
FOC (dimensionless)	0.020			İ			1		0.026744	0.025581	0.024419		0.023256	0.009884	0.012209	0.012209
SOM (calculated)	3.40%								4.61%	4.41%	4.21%		4.01%	1.70%	2.10%	2.10%
pH (su)	7.7			1			1		7.6	7.7	7.7		7.8	7.45	7.58	6.88

Risk parameter: Human health - residential with plant uptake (2.5%SOM)

Data set: MRB & WMF Client: Bovis Barratt and Taylor Wimpy Site: Land at Bankside, Banbury Job no: C12702



### Assessment of Chemicals of Potential Concern to Plant Life



								Soil Type	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT
	All values i	n mg/kg unless	otherwise	stated				Location & Depth	HTP15	HTP17	HTP20	HTP22	HTP24	HTP25	HTP26	HTP29	HTP33	HTP39	HTP40	HTP41	HTP43	HTP44
Chemical of Potential Concern	Lab. RL	No. Samples	Min. Value	Max. Value	No. Samples > or = GAC	GAC	US ₉₅	Result of Significance Test	0.25	0.15	0.15	0.20	0.20	0.15	0.10	0.15	0.25	0.20	0.20	0.1	0.2	0.7
Arsenic	2	40	17	230	0	250	130.0778	POTENTIALLY SUITABLE FOR USE	110	66	68	86	79	140	120	58	76	52	58	190	130	230
Boron	0.4	20	0.5	2.2	0	3	1.440611	POTENTIALLY SUITABLE FOR USE	0.8	0.9	1.4	0.7	0.9	0.5	1.1	1.3	1.2	0.7	0.9	1	1	0.7
Chromium (III)	5	23	46.6	589.5	3	400	329.0309	POTENTIALLY SUITABLE FOR USE	269.5	77.5	87.5	109.5	109.5	269.5	239.5	77.5	99.5	77.5	81.5	479.5	209.5	539.5
Chromium (VI)	0.5	20	0.2	1.2	0	25	0.683627	POTENTIALLY SUITABLE FOR USE	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5	0.5
Copper	5	23	6	49	0	135	38.32559	POTENTIALLY SUITABLE FOR USE	21	23	31	31	31	41	46	28	40	23	26	40	49	35
Nickel	5	40	29	210	31	75	126.9991	FURTHER ASSESSMENT REQUIRED	110	62	99	100	92	150	120	69	79	52	60	170	160	210
Zinc	10	23	74	970	5	300	393.701	FURTHER ASSESSMENT REQUIRED	190	110	180	140	160	300	260	110	150	100	120	340	290	380
	Mean																					
pH (su)	7.7								7.1	7.5	7.7	8.3	7.7	7.7	7.9	7.9	7.6	7.6	7.3	7.7	7.5	7.6
	t: MRB & V		aylor Wir	npy						-	considered	as being at	the detection	n limit for the		f statistical a	analysis, as a	dicated) and a conservati				

MG denotes Made Ground

NAT denotes natural ground

Site: Land at Bankside, Banbury

Job no: C12702

### Assessment of Chemicals of Potential Concern to Plant Life

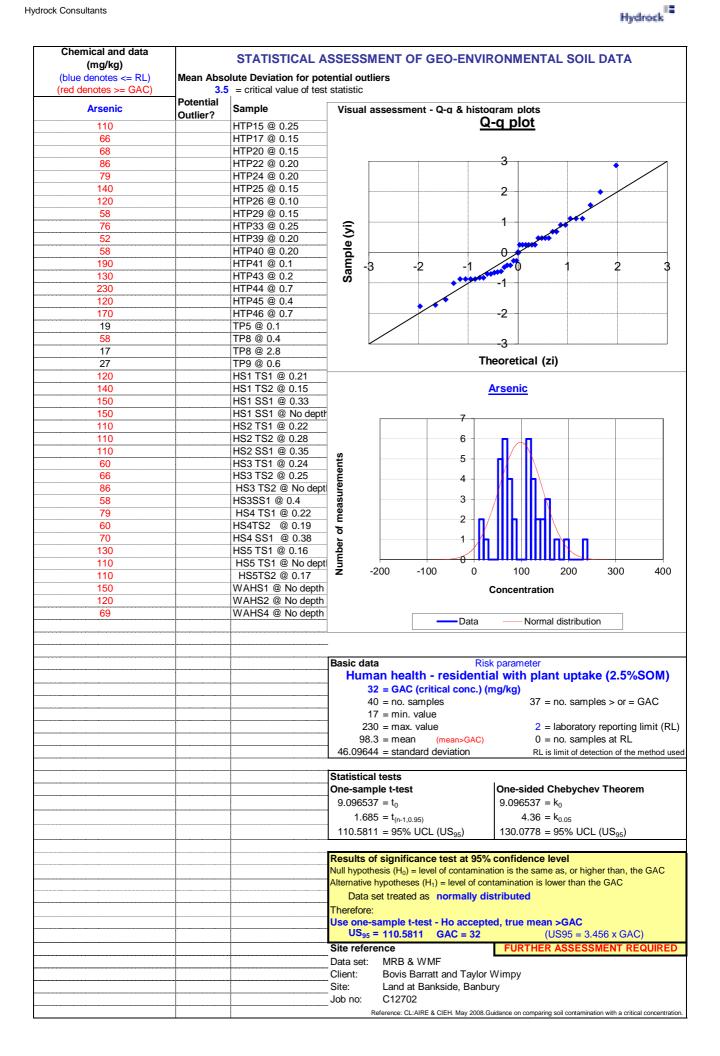


Chemical of Potential		erwise sta	ated						NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT
I an RI	No N	All values in mg/kg unless otherwise stated								TP5	TP8	TP8	TP9	HS1 TS1	HS1 TS2	HS1 SS1	HS1 SS1	HS2 TS1	HS2 TS2	HS2 SS1	HS3 TS1
Concern San		Min. /alue	Max. Value	No. Samples > or = GAC	GAC	US ₉₅	Result of Significance Test	0.4	0.7	0.1	0.4	2.8	0.6	0.21	0.15	0.33	No depth	0.22	0.28	0.35	0.24
Arsenic 2 4	40	17	230	0	250	130.0778	POTENTIALLY SUITABLE FOR USE	120	170	19	58	17	27	120	140	150	150	110	110	110	60
Boron 0.4 2	20	0.5	2.2	0	3	1.440611	POTENTIALLY SUITABLE FOR USE	1	1.2	1.8	1	2.2	0.7								
Chromium (III) 5 2	23 4	46.6	589.5	3	400	329.0309	POTENTIALLY SUITABLE FOR USE	189.5	589.5	55.57	109.8	53.8	46.6								
Chromium (VI) 0.5 2	20	0.2	1.2	0	25	0.683627	POTENTIALLY SUITABLE FOR USE	0.5	0.5	0.43	0.2	1.2	0.4								
Copper 5 2	23	6	49	0	135	38.32559	POTENTIALLY SUITABLE FOR USE	43	25	11	18	13	12								
Nickel 5 4	40	29	210	31	75	126.9991	FURTHER ASSESSMENT REQUIRED	130	180	29	57	32	43	100	100	130	150	110	110	110	75
Zinc 10 2	23	74	970	5	300	393.701	FURTHER ASSESSMENT REQUIRED	220	970	82	130	74	84								
Mean																					
pH (su) 7.7								6.9	7.9	7.2	8.1	7.4	7.1	8.2	8.1	8.1		8.1	8.1	8.1	8

### Assessment of Chemicals of Potential Concern to Plant Life

All								Soil Type	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT	NAT
1	l values in	mg/kg unless	otherwise s	stated				Location & Depth	HS3 TS2	HS3 TS2	HS3SS1	HS4 TS1	HS4TS2	HS4 SS1	HS5 TS1	HS5 TS1	HS5TS2	WAHS1	WAHS2	WAHS4
Chemical of Potential Concern	.ab. RL	No. Samples	Min. Value	Max. Value	No. Samples > or = GAC	GAC	US ₉₅	Result of Significance Test	0.25	No depth	0.4	0.22	0.19	0.38	0.16	No depth	0.17	No depth	No depth	No depti
Arsenic	2	40	17	230	0	250	130.0778	POTENTIALLY SUITABLE FOR USE	66	86	58	79	60	70	130	110	110	150	120	69
Boron	0.4	20	0.5	2.2	0	3	1.440611	POTENTIALLY SUITABLE FOR USE												
Chromium (III)	5	23	46.6	589.5	3	400	329.0309	POTENTIALLY SUITABLE FOR USE										270	150	87
Chromium (VI)	0.5	20	0.2	1.2	0	25	0.683627	POTENTIALLY SUITABLE FOR USE												
Copper	5	23	6	49	0	135	38.32559	POTENTIALLY SUITABLE FOR USE										6	6	14
Copper	5	40	29	210	31	75	126.9991	FURTHER ASSESSMENT REQUIRED	75	87	65	83	78	86	110	120	110	110	98	79
Zinc	10	23	74	970	5	300	393.701	FURTHER ASSESSMENT REQUIRED					1					310	290	170
N	Mean				1															
pH (su)	7.7								8		8	7.9	7.6	7.7	7.7		7.8	7.45	7.58	6.88

Hydrock



(mg/kg)

(blue denotes <= RL) (red denotes >= GAC)

Beryllium

4.1

1.6 2.3

2.5

2 4.2

3.7

1.6

2.1

1.4

1.6

5.4

3.6

5.6

3.3

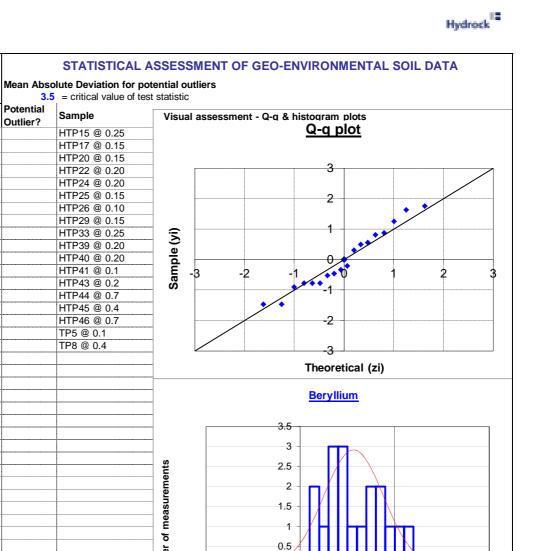
4.8

0.5

Potential

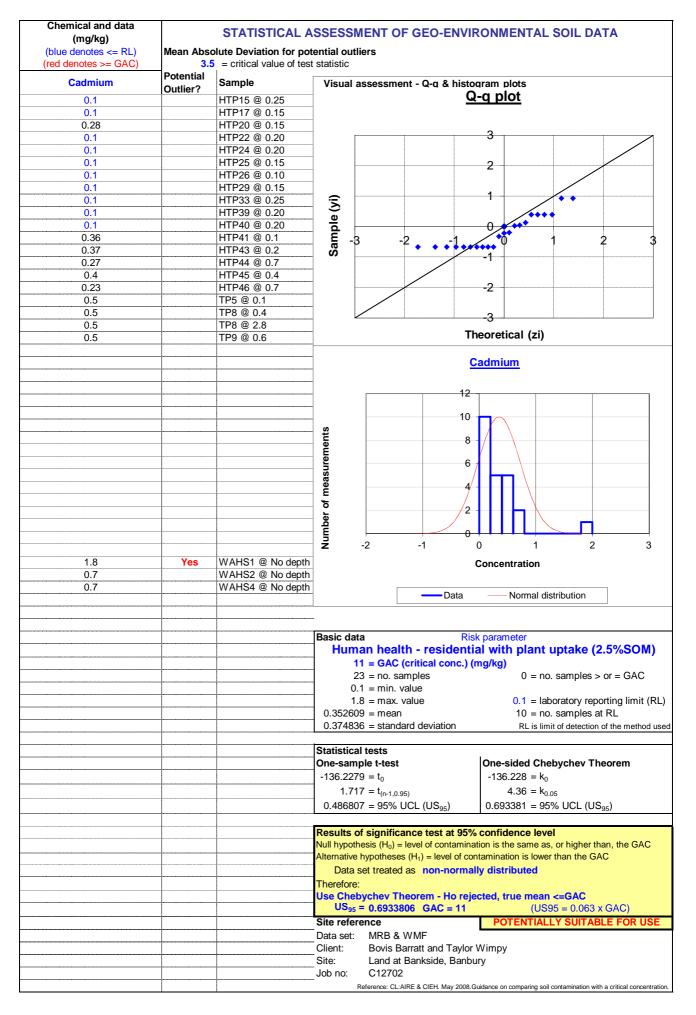
Outlier?

Sample



0.5	TP8 @ 0.4	
		Theoretical (zi)
		Beryllium
		Derymum
		3.5
		3
		2.5
		<b>i</b> 1.5
		2.5 2 1.5 1.5
		b 0.5
		-5 0 5 10
		Concentration
		Data Normal distribution
		Basic data Risk parameter
		Human health - residential with plant uptake (2.5%SOM)
		51 = GAC (critical conc.) (mg/kg)
		18 = no. samples 0 = no. samples > or = GAC
		0.5 = min. value
		5.6 = max. value 1 = laboratory reporting limit (RL)
		2.822222 = mean 2 = no. samples at RL
		1.576875 = standard deviation RL is limit of detection of the method used
		Statistical tests
		One-sample t-test One-sided Chebychev Theorem
		$-129.6241 = t_0$ $-129.624 = k_0$
		$0.174 = t_{(n-1,0.95)}                                     $
		$2.886893 = 95\% \text{ UCL } (\text{US}_{95}) \qquad 4.442717 = 95\% \text{ UCL } (\text{US}_{95})$
		$2.000033 - 35\% \text{ OCL } (0395) \qquad 4.442717 = 35\% \text{ OCL } (0395)$
		Results of significance test at 95% confidence level
		Null hypothesis ( $H_0$ ) = level of contamination is the same as, or higher than, the GAC
		Alternative hypotheses ( $H_1$ ) = level of contamination is lower than the GAC
		Data set treated as non-normally distributed
		Therefore:
		Use Chebychev Theorem - Ho rejected, true mean <=GAC
		US ₉₅ = 4.4427169 GAC = 51 (US95 = 0.087 x GAC)
		Site reference POTENTIALLY SUITABLE FOR USE
		Data set: MRB & WMF
		Client: Bovis Barratt and Taylor Wimpy
		Site: Land at Bankside, Banbury
		Job no: C12702
		Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.

Chemical and data (mg/kg)		STATISTICAL	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL)	Mean Abso	olute Deviation for po	otential outliers
(red denotes >= GAC)		= critical value of tes	
Boron	Potential	Sample	Visual assessment - Q-q & histogram plots
0.8	Outlier?	HTP15 @ 0.25	Q-q plot
0.9		HTP17 @ 0.15	
1.4		HTP20 @ 0.15	-
0.7		HTP22 @ 0.20	3
0.9		HTP24 @ 0.20	
0.5		HTP25 @ 0.15	2
1.1		HTP26 @ 0.10	
1.3		HTP29 @ 0.15	
1.2		HTP33 @ 0.25	Sample (y)
0.7		HTP39 @ 0.20 HTP40 @ 0.20	
1		HTP41 @ 0.1	
1		HTP43 @ 0.2	
0.7		HTP44 @ 0.7	1
1		HTP45 @ 0.4	
1.2		HTP46 @ 0.7	-2
1.8		TP5 @ 0.1	
1		TP8 @ 0.4	
2.2	Yes	TP8 @ 2.8	Theoretical (zi)
0.7		TP9 @ 0.6	
			Boron
			_
			5
			Number of measurements
		_	<b>ž</b> -1 0 1 2 3
			Concentration
			Data — Normal distribution
			www.
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			290 = GAC (critical conc.) (mg/kg)
			20 = no.  samples $0 = no.  samples > or = GAC$
			0.5 = min. value
			2.2 = max. value 0.4 = laboratory reporting limit (RL)
			1.05 = mean 0 = no. samples at RL
			0.400657 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-3225.259 = t_0$ $-3225.26 = k_0$
			$1.729 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			$1.204901 = 95\% \text{ UCL } (\text{US}_{95})$ $1.440611 = 95\% \text{ UCL } (\text{US}_{95})$
			Results of significance test at 95% confidence level
			Null hypothesis $(H_0)$ = level of contamination is the same as, or higher than, the GAC Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
			Data set treated as <b>non-normally distributed</b>
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			$US_{95} = 1.4406111$ GAC = 290 (US95 = 0.005 x GAC)
	1	1	
			Site reference POTENTIALLY SUITABLE FOR USE
			Site reference         POTENTIALLY SUITABLE FOR USE           Data set:         MRB & WMF
			Data set: MRB & WMF
			Data set: MRB & WMF Client: Bovis Barratt and Taylor Wimpy





Γ

Chemical and data

(blue denotes <= RL)		STATISTICAL A	SSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(red denotes >= GAC)		<b>Dilute Deviation for pot</b> = critical value of test	
Chromium (III)	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots
269.5	Cullor	HTP15 @ 0.25	<u>Q-q plot</u>
77.5		HTP17 @ 0.15	
87.5		HTP20 @ 0.15	
109.5		HTP22 @ 0.20	3
109.5		HTP24 @ 0.20	
269.5		HTP25 @ 0.15	2
239.5		HTP26 @ 0.10	
77.5		HTP29 @ 0.15	
99.5		HTP33 @ 0.25	
77.5		HTP39 @ 0.20	
81.5		HTP40 @ 0.20	
479.5	Yes	HTP41 @ 0.1	
209.5		HTP43 @ 0.2	Sample ( <i>X</i> i)
539.5	Yes	HTP44 @ 0.7	
189.5		HTP45 @ 0.4	
589.5	Yes	HTP46 @ 0.7	-2
55.57		TP5 @ 0.1	
109.8		TP8 @ 0.4	-3
53.8		TP8 @ 2.8	-
46.6		TP9 @ 0.6	Theoretical (zi)
			Chromium (III)
			10
			9 9
			5 - 1
			გ. 2 - 1
			Number of the second se
270		WAHS1 @ No depth	-1000 -500 0 500 1000 Concentration
150		WAHS2 @ No depth	
150		WAHS2 @ No depth	Concentration
150		WAHS2 @ No depth	Concentration Data Normal distribution
150		WAHS2 @ No depth	Concentration Data Normal distribution Basic data Risk parameter
150		WAHS2 @ No depth	Concentration — Data — Normal distribution Basic data Risk parameter Human health - residential with plant uptake (2.5%SOM)
150		WAHS2 @ No depth	Concentration — Data Normal distribution Basic data Risk parameter Human health - residential with plant uptake (2.5%SOM) 630 = GAC (critical conc.) (mg/kg)
150		WAHS2 @ No depth	Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)         630 = GAC (critical conc.) (mg/kg)         23 = no. samples       0 = no. samples > or = GAC
150		WAHS2 @ No depth	Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)         630 = GAC (critical conc.) (mg/kg)         23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value
150		WAHS2 @ No depth	Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)       630 = GAC (critical conc.) (mg/kg)         23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value       5 = laboratory reporting limit (RL)
150		WAHS2 @ No depth	Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)       630 = GAC (critical conc.) (mg/kg)         23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value       5 = laboratory reporting limit (RL)         589.5 = max. value       5 = laboratory reporting limit (RL)         186.0335 = mean       0 = no. samples at RL
150		WAHS2 @ No depth	Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)       630 = GAC (critical conc.) (mg/kg)         23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value       5 = laboratory reporting limit (RL)         589.5 = max. value       5 = no. samples at RL
150		WAHS2 @ No depth	Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)       630 = GAC (critical conc.) (mg/kg)         23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value       5 = laboratory reporting limit (RL)         589.5 = max. value       5 = no. samples at RL
150		WAHS2 @ No depth	Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)       630 = GAC (critical conc.) (mg/kg)         23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value       5 = laboratory reporting limit (RL)         589.5 = max. value       5 = laboratory reporting limit (RL)         186.0335 = mean       0 = no. samples at RL         157.2917 = standard deviation       RL is limit of detection of the method used
150		WAHS2 @ No depth	Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)       630 = GAC (critical conc.) (mg/kg)         23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value       5 = laboratory reporting limit (RL)         589.5 = max. value       5 = laboratory reporting limit (RL)         186.0335 = mean       0 = no. samples at RL         157.2917 = standard deviation       RL is limit of detection of the method used         Statistical tests
150		WAHS2 @ No depth	Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)       630 = GAC (critical conc.) (mg/kg)         23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value       5 = laboratory reporting limit (RL)         55 = max. value       5 = laboratory reporting limit (RL)         186.0335 = mean       0 = no. samples at RL         157.2917 = standard deviation       RL is limit of detection of the method use         Statistical tests       One-sided Chebychev Theorem         -13.53665 = t ₀ -13.5366 = k ₀
150		WAHS2 @ No depth	Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)       630 = GAC (critical conc.) (mg/kg)         23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value       5 = laboratory reporting limit (RL)         589.5 = max. value       5 = laboratory reporting limit (RL)         186.0335 = mean       0 = no. samples at RL         157.2917 = standard deviation       RL is limit of detection of the method user         Statistical tests         One-sided Chebychev Theorem         -13.53665 = t ₀ -13.5366 = k ₀ 1.717 = t _(n-1,0.95) 4.36 = k _{0.05}
150		WAHS2 @ No depth	Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)       630 = GAC (critical conc.) (mg/kg)         23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value       5 = laboratory reporting limit (RL)         55 = max. value       5 = laboratory reporting limit (RL)         186.0335 = mean       0 = no. samples at RL         157.2917 = standard deviation       RL is limit of detection of the method used         Statistical tests       One-sided Chebychev Theorem         -13.53665 = t ₀ -13.5366 = k ₀
150		WAHS2 @ No depth	Concentration         Concentration         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)       630 = GAC (critical conc.) (mg/kg)         23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value       5 = laboratory reporting limit (RL)         589.5 = max. value       5 = laboratory reporting limit (RL)         186.0335 = mean       0 = no. samples at RL         157.2917 = standard deviation       RL is limit of detection of the method used         Statistical tests         One-sided Chebychev Theorem         -13.53665 = t ₀ -13.5366 = k ₀ 1.717 = t _(n-1,0.95) 4.36 = k _{0.05}
150		WAHS2 @ No depth	ConcentrationConcentrationDataNormal distributionBasic dataRisk parameterHuman health - residential with plant uptake (2.5%SOM)630 = GAC (critical conc.) (mg/kg)23 = no. samples0 = no. samples > or = GAC46.6 = min. value5 = laboratory reporting limit (RL)589.5 = max. value5 = laboratory reporting limit (RL)186.0335 = mean0 = no. samples at RL157.2917 = standard deviationRL is limit of detection of the method usedStatistical tests One-sample t-test -13.53665 = t_0 1.717 = t_{(n-1,0.95)} 242.3469 = 95% UCL (US_{95})One-sided Chebychev Theorem-13.5366 = k_0 4.36 = k_{0.05} 329.0309 = 95% UCL (US_{95})Results of significance test at 95% confidence levelNull hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
150		WAHS2 @ No depth	Concentration         Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)         630 = GAC (critical conc.) (mg/kg) $23 = no. samples$ $0 = no. samples > or = GAC$ 46.6 = min. value $5 = laboratory reporting limit (RL)$ 55 = max. value $5 = laboratory reporting limit (RL)$ 186.0335 = mean $0 = no. samples at RL$ 157.2917 = standard deviation       RL is limit of detection of the method used         Statistical tests         One-sided Chebychev Theorem         -13.53665 = t ₀ $-13.53666 = k_0$ $1.717 = t_{(n-10.95)}$ $242.3469 = 95\%$ UCL (US ₉₅ )         Statistical tests         One-sided Chebychev Theorem         -13.53666 = k_0 $4.36 = k_{0.05}$ $242.3469 = 95\%$ UCL (US ₉₅ ) $329.0309 = 95\%$ UCL (US ₉₅ )
150		WAHS2 @ No depth	ConcentrationConcentrationDataNormal distributionBasic dataRisk parameterHuman health - residential with plant uptake (2.5%SOM)630 = GAC (critical conc.) (mg/kg)23 = no. samples0 = no. samples > or = GAC46.6 = min. value5 = laboratory reporting limit (RL)589.5 = max. value5 = laboratory reporting limit (RL)186.0335 = mean0 = no. samples at RL157.2917 = standard deviationRL is limit of detection of the method usedStatistical testsOne-sided Chebychev Theorem-13.53665 = t_0-13.5366 = k_01.717 = t _(n-1,0.95) 242.3469 = 95% UCL (US ₉₅ )Results of significance test at 95% confidence levelNull hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
150		WAHS2 @ No depth	Concentration         Concentration         Data       Normal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)         630 = GAC (critical conc.) (mg/kg)       23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value       5 = laboratory reporting limit (RL)         589.5 = max. value       5 = laboratory reporting limit (RL)         186.0335 = mean       0 = no. samples at RL         157.2917 = standard deviation       RL is limit of detection of the method used         Statistical tests         One-sided Chebychev Theorem         -13.53665 = t_0       -13.5366 = k_0         1.717 = t _(n-1,0.95) 242.3469 = 95% UCL (US ₉₅ )         Results of significance test at 95% confidence level         Null hypothesis (H ₀ ) = level of contamination is the same as, or higher than, the GAC         Alternative hypotheses (H ₁ ) = level of contamination is lower than the GAC
150		WAHS2 @ No depth	ConcentrationConcentrationDataNormal distributionBasic dataRisk parameterHuman health - residential with plant uptake (2.5%SOM)630 = GAC (critical conc.) (mg/kg)23 = no. samples0 = no. samples > or = GAC46.6 = min. value5 = laboratory reporting limit (RL)589.5 = max. value5 = laboratory reporting limit (RL)186.0335 = mean0 = no. samples at RL157.2917 = standard deviationRL is limit of detection of the method usedStatistical testsOne-sided Chebychev Theorem-13.53666 = t_0-13.5366 = k_01.717 = t _(n-1,0.95) 242.3469 = 95% UCL (US ₉₅ )Results of significance test at 95% confidence levelNull hypothesis (H_0) = level of contamination is the same as, or higher than, the GACAlternative hypotheses (H_1) = level of contamination is lower than the GACAlternative hypotheses (H_1) = level of contamination is lower than the GACData set treated as non-normally distributed
150		WAHS2 @ No depth	Concentration         Concentration         Concentration         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)         630 = GAC (critical conc.) (mg/kg)       23 = no. samples       0 = no. samples > or = GAC         46.6 = min. value       5 = laboratory reporting limit (RL)         589.5 = max. value       5 = laboratory reporting limit (RL)         186.0335 = mean       0 = no. samples at RL         157.2917 = standard deviation       RL is limit of detection of the method used         Statistical tests         One-sided Chebychev Theorem         -13.53665 = t_0       -13.5366 = k_0         1.717 = t _(n-1,0.95) 242.3469 = 95% UCL (US ₉₅ )         Zesults of significance test at 95% confidence level         Null hypothesis (H ₀ ) = level of contamination is the same as, or higher than, the GAC         Alternative hypotheses (H ₁ ) = level of contamination is lower than the GAC         Data set treated as non-normally distributed         Therefore:
150		WAHS2 @ No depth	ConcentrationConcentration— Data Normal distributionBasic data Risk parameterHuman health - residential with plant uptake (2.5%SOM) $630 = GAC$ (critical conc.) (mg/kg) $23 = no.$ samples $0 = no.$ samples > or = GAC $46.6 = min.$ value $5 =$ laboratory reporting limit (RL) $589.5 = max.$ value $5 =$ laboratory reporting limit (RL) $186.0335 =$ mean $0 = no.$ samples at RL $157.2917 =$ standard deviationRL is limit of detection of the method usedStatistical testsOne-sided Chebychev Theorem $-13.53656 = t_0$ $-13.5366 = k_0$ $1.717 = t_{(n-1,0.95)}$ $242.3469 = 95\%$ UCL (US ₉₅ )Results of significance test at 95% confidence levelNull hypothesis (H_0) = level of contamination is the same as, or higher than, the GACAlternative hypotheses (H_1) = level of contamination is lower than the GACData set treated as non-normally distributedTherefore:Use Chebychev Theorem - Ho rejected, true mean <=GAC
150		WAHS2 @ No depth	ConcentrationConcentrationConcentrationDataNormal distributionBasic dataRisk parameterHuman health - residential with plant uptake (2.5% SOM) $630 = GAC$ (critical conc.) (mg/kg) $23 = no.$ samples $0 = no.$ samples > or = GAC $46.6 = min.$ value $5 = laboratory reporting limit (RL)589.5 = max. value5 = laboratory reporting limit (RL)186.0335 = mean0 = no. samples at RL157.2917 = standard deviationRL is limit of detection of the method userStatistical testsOne-sided Chebychev Theorem-13.53656 = t_0-13.5366 = k_01.717 = t_{(n-1,0.95)}329.0309 = 95\% UCL (US95)Results of significance test at 95% confidence levelNull hypothesis (H0) = level of contamination is the same as, or higher than, the GACAlternative hypotheses (H1) = level of contamination is lower than the GACData set treated as non-normally distributedTherefore:Use Chebychev Theorem - Ho rejected, true mean <=GACUS95 = 329.03091GAC = 630(US95 = 0.522 x GAC)$
150		WAHS2 @ No depth	ConcentrationConcentration <b>Basic data</b> Risk parameterHuman health - residential with plant uptake (2.5%SOM)630 = GAC (critical conc.) (mg/kg)23 = no. samples0 = no. samples > or = GAC46.6 = min. value5 = laboratory reporting limit (RL)589.5 = max. value5 = laboratory reporting limit (RL)186.0335 = mean0 = no. samples at RL157.2917 = standard deviationRL is limit of detection of the method userStatistical testsOne-sided Chebychev Theorem-13.53656 = to-13.5366 = k_01.717 = t _(n-1,0.95) 242.3469 = 95% UCL (US ₉₅ )Z42.3469 = 95% UCL (US ₉₅ )Results of significance test at 95% confidence levelNull hypothesis (H ₀ ) = level of contamination is the same as, or higher than, the GACAlternative hypotheses (H ₁ ) = level of contamination is lower than the GACData set treated as non-normally distributedTherefore:Use Chebychev Theorem - Ho rejected, true mean <=GACUS95 = 329.03091 GAC = 630(US95 = 0.522 × GAC)Site referencePOTENTIALLY SUITABLE FOR USE
150		WAHS2 @ No depth	Concentration         Concentration         Mormal distribution         Basic data       Risk parameter         Human health - residential with plant uptake (2.5%SOM)         630 = GAC (critical conc.) (mg/kg) $23 = no. samples$ $0 = no. samples > or = GAC$ 46.6 = min. value $5 =$ laboratory reporting limit (RL)         186.0335 = mean $0 = no. samples at RL$ 157.2917 = standard deviation       RL is limit of detection of the method user         Statistical tests         One-sided Chebychev Theorem         -13.53656 = to       1.717 = t _(n-1,0.95) 242.3469 = 95% UCL (US ₉₅ )       329.0309 = 95% UCL (US ₉₅ )         Results of significance test at 95% confidence level         Null hypothesis (H ₀ ) = level of contamination is the same as, or higher than, the GAC         Alternative hypotheses (H ₁ ) = level of contamination is lower than the GAC         Alternative hypotheses (H ₁ ) = level of contamination is lower than the GAC         Use Chebychev Theorem - Ho rejected, true mean <=GAC         US ₉₅ = 329.03091       GAC = 630       (US95 = 0.522 × GAC)         Site reference         POTENTIALLY SUITABLE FOR USE         Data set:       MRB & WMF
150		WAHS2 @ No depth	ConcentrationConcentration <b>Basic data</b> Risk parameterHuman health - residential with plant uptake (2.5%SOM)630 = GAC (critical conc.) (mg/kg)23 = no. samples0 = no. samples > or = GAC46.6 = min. value5 = laboratory reporting limit (RL)589.5 = max. value5 = laboratory reporting limit (RL)186.0335 = mean0 = no. samples at RL157.2917 = standard deviationRL is limit of detection of the method userStatistical testsOne-sided Chebychev Theorem-13.53656 = to-13.5366 = k_01.717 = t _(n-1,0.95) 242.3469 = 95% UCL (US ₉₅ )Z42.3469 = 95% UCL (US ₉₅ )Results of significance test at 95% confidence levelNull hypothesis (H ₀ ) = level of contamination is the same as, or higher than, the GACAlternative hypotheses (H ₁ ) = level of contamination is lower than the GACData set treated as non-normally distributedTherefore:Use Chebychev Theorem - Ho rejected, true mean <=GACUS95 = 329.03091 GAC = 630(US95 = 0.522 × GAC)Site referencePOTENTIALLY SUITABLE FOR USE

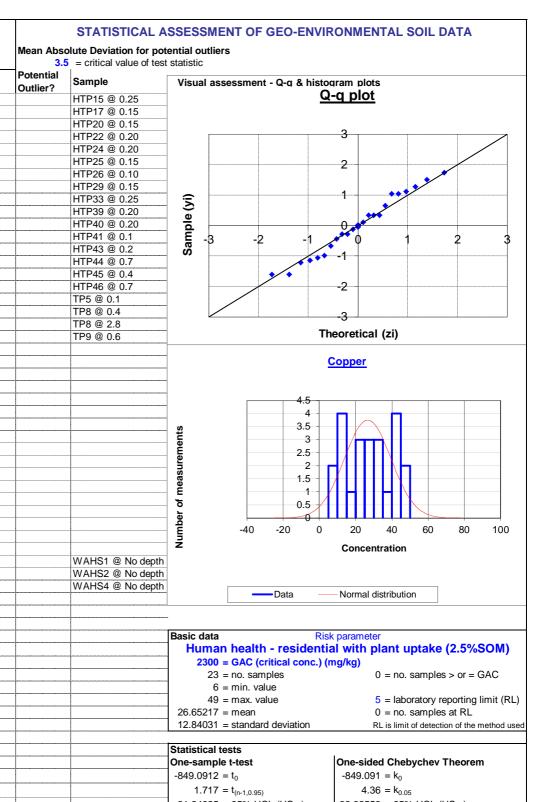
Chemical and data (mg/kg) (blue denotes <= RL) (red denotes >= GAC)		STATISTICAL olute Deviation for p = critical value of to	
	Potential		•••
Chromium (VI)	Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.5	n/a	HTP15 @ 0.25	<u>Q-q plot</u>
0.5	n/a	HTP17 @ 0.15	
0.5	n/a	HTP20 @ 0.15	3
0.5	n/a	HTP22 @ 0.20	
0.5	n/a	HTP24 @ 0.20	
0.5	n/a n/a	HTP25 @ 0.15 HTP26 @ 0.10	2
0.5	n/a	HTP29 @ 0.15	
0.5	n/a	HTP33 @ 0.25	
0.5	n/a	HTP39 @ 0.20	
0.5	n/a	HTP40 @ 0.20	
0.5	n/a	HTP41 @ 0.1	Sample (yi)
0.5	n/a	HTP43 @ 0.2	
0.5	n/a	HTP44 @ 0.7	
0.5	n/a	HTP45 @ 0.4	
0.5	n/a	HTP46 @ 0.7	-2 -2
0.43	n/a n/a	TP5 @ 0.1 TP8 @ 0.4	
1.2	n/a	TP8 @ 2.8	
0.4	n/a	TP9 @ 0.6	Theoretical (zi)
			Chromium (VI)
			_
			o 12
			─┤ ┋
			<b>ž</b> -0.5 0 0.5 1 1.5
			Concentration
			Data Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			4.3 = GAC (critical conc.) (mg/kg)
		1	20 = no. samples $0 = no. samples > or = GAC$
			0.2 = min. value
			1.2 = max. value 0.5 = laboratory reporting limit (RL)
			0.5115 = mean 19 = no. samples at RL
			0.176554 = standard deviation RL is limit of detection of the method used
			0(-0
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-95.9632 = t_0$ $-95.9632 = k_0$
			$1.729 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			0.579759 = 95% UCL (US ₉₅ ) $0.683627 = 95%$ UCL (US ₉₅ )
			Desults of significance tost at 05% confidence level
			<b>Results of significance test at 95% confidence level</b> Null hypothesis ( $H_0$ ) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
			Data set treated as <b>non-normally distributed</b>
		1	Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			$US_{95} = 0.683627$ GAC = 4.3 (US95 = 0.159 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: MRB & WMF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.

(mg/kg)

(blue denotes <= RL)

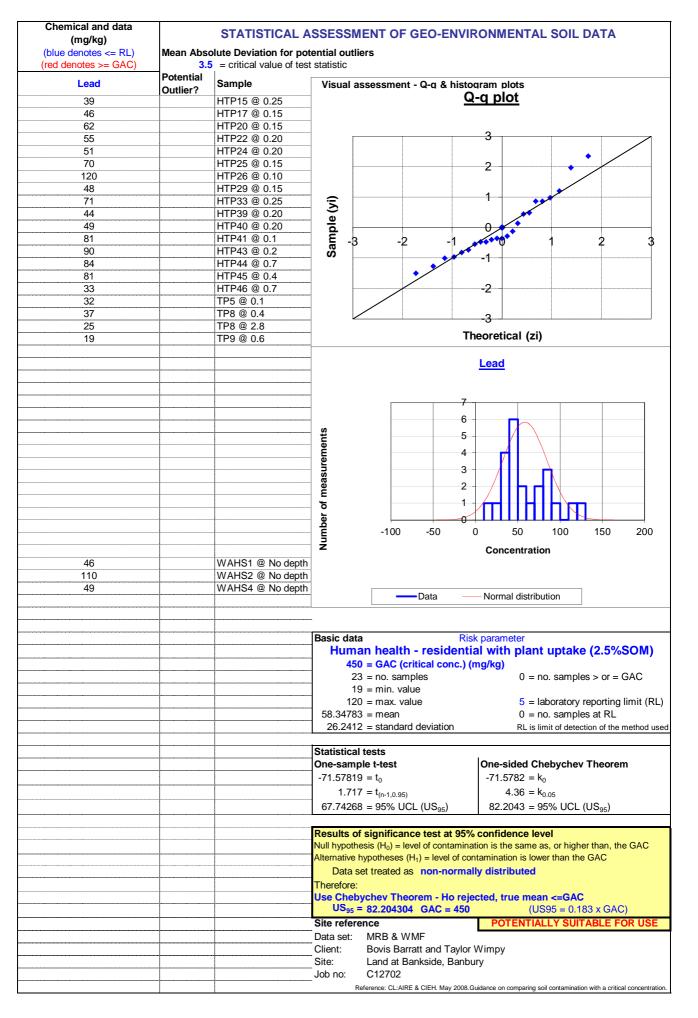
(red denotes >= GAC)

Copper

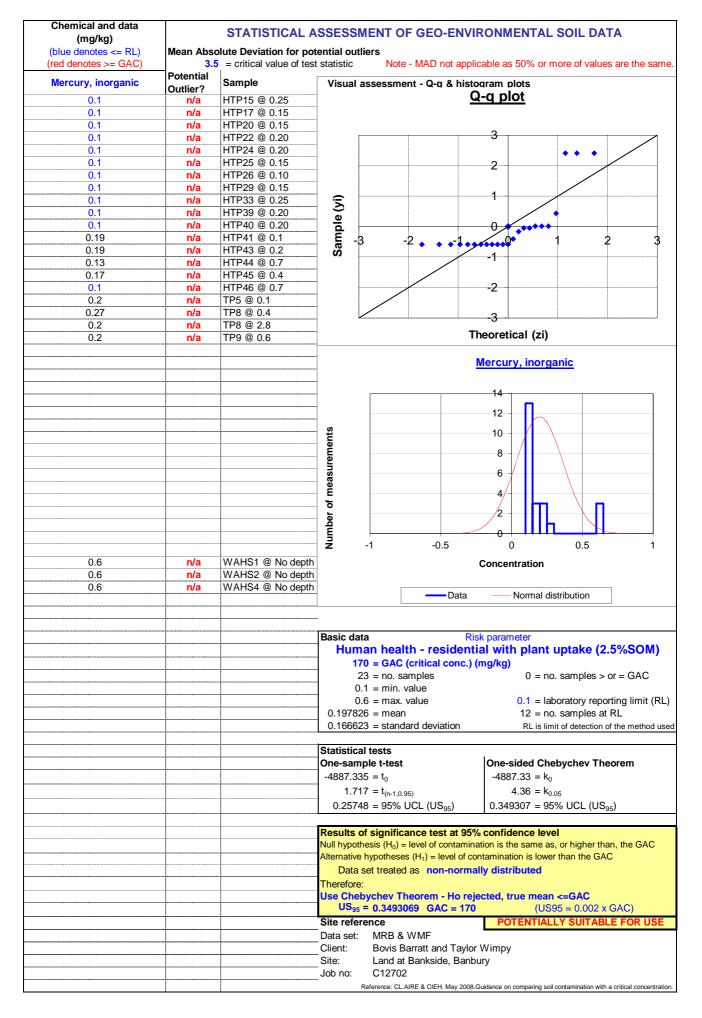


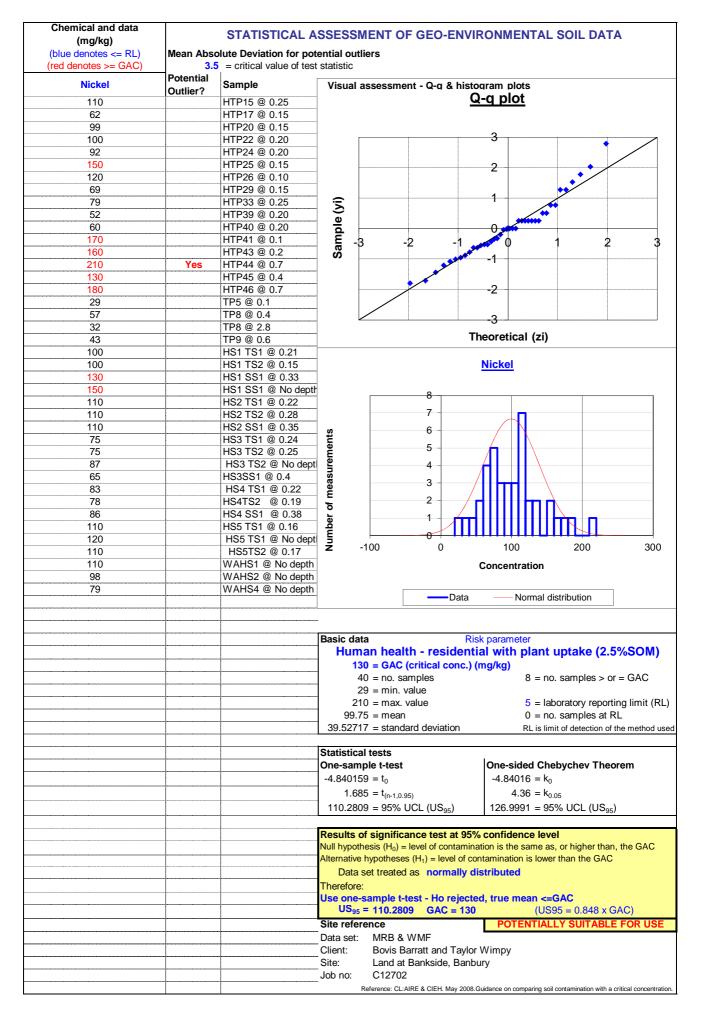
	On a side d Ob should be The second
One-sample t-test	One-sided Chebychev Theorem
-849.0912 = t ₀	$-849.091 = k_0$
$1.717 = t_{(n-1,0.95)}$	$4.36 = k_{0.05}$
31.24925 = 95% UCL (US ₉₅ )	38.32559 = 95% UCL (US ₉₅ )
Results of significance test at	95% confidence level
Null hypothesis $(H_0) =$ level of containing the second	mination is the same as, or higher than, the GAC
Alternative hypotheses (H ₁ ) = level of	of contamination is lower than the GAC
Data set treated as non-nor	rmally distributed
Therefore:	
Use Chebychev Theorem - Ho	rejected, true mean <=GAC
US ₉₅ = 38.325592 GAC = 2	
Site reference	POTENTIALLY SUITABLE FOR USE
Data set: MRB & WMF	
Client: Bovis Barratt and Ta	ylor Wimpy
Site: Land at Bankside, Ba	anbury
Job no: C12702	
Reference: CL:AIRE & CIEH. May 2	2008. Guidance on comparing soil contamination with a critical concentration.

Hydrock



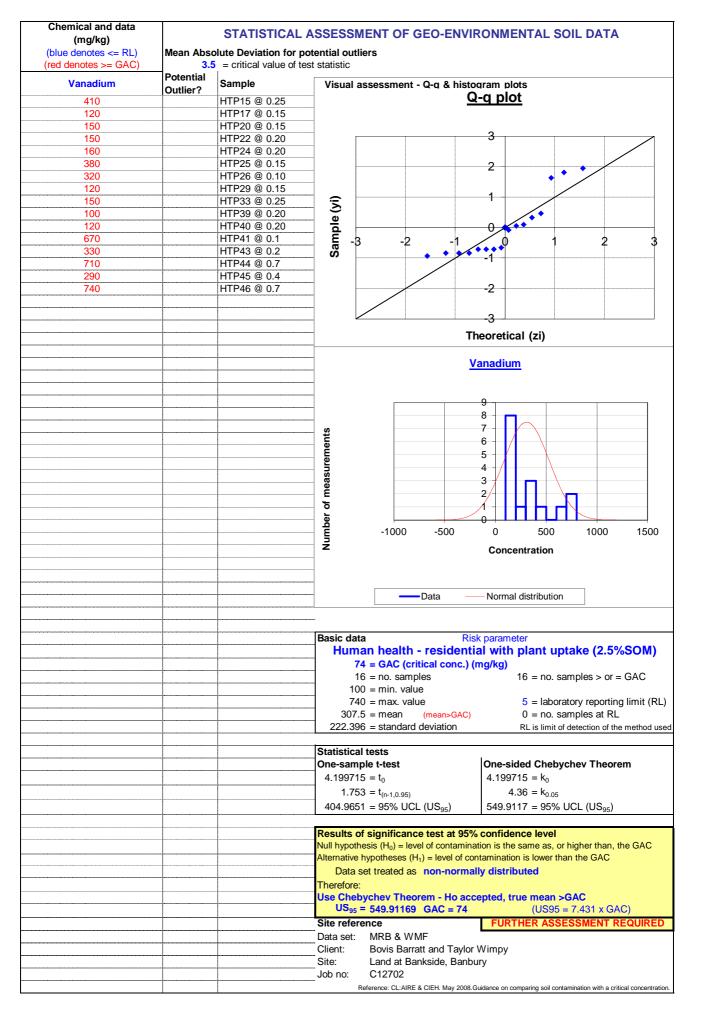




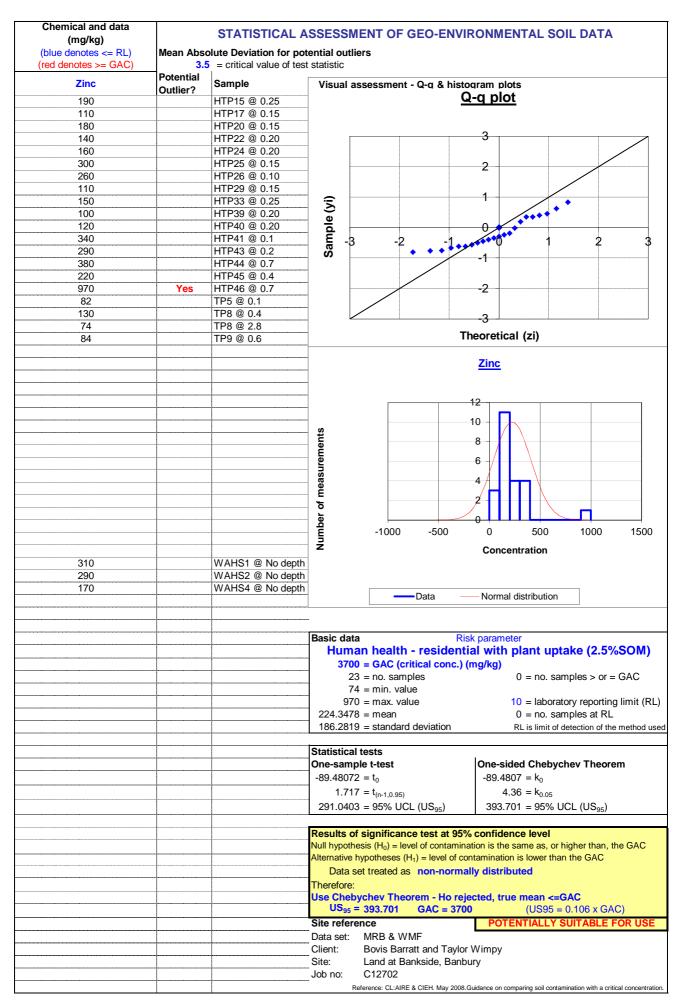




Chemical and data (mg/kg) (blue denotes <= RL) (red denotes >= GAC)		STATISTICAL A blute Deviation for pot 5 = critical value of test	ential outlier	
(red denotes >= GAC)			statistic	Note - MAD not applicable as 50% or more of values are the same.
Selenium	Outlier?	Sample	Visual ass	essment - Q-q & histogram plots
 0.2	n/a	HTP15 @ 0.25		<u>Q-q plot</u>
0.2	n/a	HTP17 @ 0.15		
0.2	n/a	HTP20 @ 0.15		
0.2	n/a	HTP22 @ 0.20		3
 0.2	n/a	HTP24 @ 0.20		•••
 0.22	n/a	HTP25 @ 0.15		2
 0.2	n/a	HTP26 @ 0.10		
 0.2	n/a	HTP29 @ 0.15	<b>~</b>	1
 0.2	n/a n/a	HTP33 @ 0.25 HTP39 @ 0.20	Sample (yi)	
 0.2	n/a	HTP40 @ 0.20	e	
 0.2	n/a	HTP41 @ 0.1	du	-2 • • • • • • • • • • • • • • • • • • •
 0.2	n/a	HTP43 @ 0.2	β-B	-2 - 2 - 3
 0.2	n/a	HTP44 @ 0.7	0,	-1
 0.2	n/a	HTP45 @ 0.4		
 0.2	n/a	HTP46 @ 0.7		-2
 0.3	n/a	TP5 @ 0.1		
 0.3	n/a	TP8 @ 0.4		
 0.3	n/a	TP8 @ 2.8		6
 0.3	n/a	TP9 @ 0.6		Theoretical (zi)
 				<u>Selenium</u>
 			irements	20
			Number of measurements	
 			ž	Concentration
 3	n/a	WAHS1 @ No depth		
3	n/a	WAHS2 @ No depth		
 3	n/a	WAHS4 @ No depth		
				Data      Normal distribution
 			Basic data	Risk parameter
 				health - residential with plant uptake (2.5%SOM)
 				= GAC (critical conc.) (mg/kg)
				= no. samples 0 = no. samples > or = GAC
		1	0.2	
 			-	= min. value
 			3 =	= max. value 0.2 = laboratory reporting limit (RL)
 			3 = 0.583478 =	= max. value0.2 = laboratory reporting limit (RL)= mean15 = no. samples at RL
 			3 = 0.583478 =	= max. value0.2 = laboratory reporting limit (RL)= mean15 = no. samples at RL
			3 = 0.583478 = 0.957699 =	= max. value       0.2 = laboratory reporting limit (RL)         = mean       15 = no. samples at RL         = standard deviation       RL is limit of detection of the method used
			3 = 0.583478 = 0.957699 = Statistical te	= max. value     0.2 = laboratory reporting limit (RL)       = mean     15 = no. samples at RL       = standard deviation     RL is limit of detection of the method used
			3 = 0.583478 = 0.957699 = Statistical te One-sample	= max. value     0.2 = laboratory reporting limit (RL)       = mean     15 = no. samples at RL       = standard deviation     RL is limit of detection of the method used       ests     one-sided Chebychev Theorem
			3 = 0.583478 = 0.957699 = Statistical te One-sample -1749.759 =	= max. value     0.2 = laboratory reporting limit (RL)       = mean     15 = no. samples at RL       = standard deviation     RL is limit of detection of the method used       ests     • t-test       = t_0     -1749.76 = k_0
			3 = 0.583478 = 0.957699 = Statistical te One-sample -1749.759 = 1.717 =	= max. value         0.2 = laboratory reporting limit (RL)           = mean         15 = no. samples at RL           = standard deviation         RL is limit of detection of the method used           ests         • t-test           = t ₀ -1749.76 = k ₀ = t _(n-1,0.95) 4.36 = k _{0.05}
			3 = 0.583478 = 0.957699 = Statistical te One-sample -1749.759 = 1.717 =	= max. value     0.2 = laboratory reporting limit (RL)       = mean     15 = no. samples at RL       = standard deviation     RL is limit of detection of the method used       ests     • t-test       = t_0     -1749.76 = k_0
			3 = 0.583478 = 0.957699 = <b>Statistical te</b> <b>One-sample</b> -1749.759 = 1.717 = 0.926353 =	= max. value $0.2$ = laboratory reporting limit (RL)= mean15 = no. samples at RL= standard deviationRL is limit of detection of the method usedestsOne-sided Chebychev Theorem= $t_0$ -1749.76 = $k_0$ = $t_{(n-1,0.95)}$ 4.36 = $k_{0.05}$ = 95% UCL (US ₉₅ )1.454144 = 95% UCL (US ₉₅ )
			3 = 0.583478 = 0.957699 = <b>Statistical te</b> <b>One-sample</b> -1749.759 = 1.717 = 0.926353 = <b>Results of s</b>	= max. value $0.2$ = laboratory reporting limit (RL)= mean $15$ = no. samples at RL= standard deviationRL is limit of detection of the method usedestsOne-sided Chebychev Theorem= $t_0$ $-1749.76 = k_0$ = $t_{(n-1,0.95)}$ $4.36 = k_{0.05}$ = 95% UCL (US ₉₅ ) $1.454144 = 95\%$ UCL (US ₉₅ )
			3 = 0.583478 = 0.957699 = <b>Statistical te</b> <b>One-sample</b> -1749.759 = 1.717 = 0.926353 = <b>Results of s</b> Null hypothesi	= max. value $0.2$ = laboratory reporting limit (RL)         = mean $15$ = no. samples at RL         = standard deviation       RL is limit of detection of the method used         ests $1749.76 = k_0$ = t _(n-1,0.95) $4.36 = k_{0.05}$ = 95% UCL (US ₉₅ ) $1.454144 = 95\%$ UCL (US ₉₅ )         ignificance test at 95% confidence level         is (H ₀ ) = level of contamination is the same as, or higher than, the GAC
			3 = 0.583478 = 0.957699 = <b>Statistical te</b> <b>One-sample</b> -1749.759 = 1.717 = 0.926353 = <b>Results of s</b> Null hypothes Alternative hyp	= max. value       0.2 = laboratory reporting limit (RL)         = mean       15 = no. samples at RL         = standard deviation       RL is limit of detection of the method used         ests       • t-test         = $t_0$ -1749.76 = $k_0$ = $t_{(n-1,0.95)}$ 4.36 = $k_{0.05}$ = 95% UCL (US ₉₅ )       1.454144 = 95% UCL (US ₉₅ )         ignificance test at 95% confidence level         is (H_0) = level of contamination is the same as, or higher than, the GAC potheses (H_1) = level of contamination is lower than the GAC
			3 = 0.583478 = 0.957699 = <b>Statistical te</b> <b>One-sample</b> -1749.759 = 1.717 = 0.926353 = <b>Results of s</b> Null hypothes Alternative hypothes Data set	= max. value $0.2$ = laboratory reporting limit (RL)         = mean $15$ = no. samples at RL         = standard deviation       RL is limit of detection of the method used         ests $1749.76 = k_0$ = t _(n-1,0.95) $4.36 = k_{0.05}$ = 95% UCL (US ₉₅ ) $1.454144 = 95\%$ UCL (US ₉₅ )         ignificance test at 95% confidence level         is (H ₀ ) = level of contamination is the same as, or higher than, the GAC
			3 = 0.583478 = 0.957699 = <b>Statistical te</b> <b>One-sample</b> -1749.759 = 1.717 = 0.926353 = <b>Results of s</b> Null hypothesi Alternative hypothesi Data set Therefore:	= max. value       0.2 = laboratory reporting limit (RL)         = mean       15 = no. samples at RL         = standard deviation       RL is limit of detection of the method used         ests       • t-test         = t_{0}       -1749.76 = k_0         = t_{(n-1,0.95)}       4.36 = k_{0.05}         = 95% UCL (US ₉₅ )       1.454144 = 95% UCL (US ₉₅ )         ignificance test at 95% confidence level       is (H_0) = level of contamination is the same as, or higher than, the GAC potheses (H_1) = level of contamination is lower than the GAC treated as non-normally distributed
			3 = 0.583478 = 0.957699 = <b>Statistical te</b> <b>One-sample</b> -1749.759 = 1.717 = 0.926353 = <b>Results of s</b> Null hypothesi Alternative hypothesi Alternative hypothesi Data set Therefore: Use Chebyo	= max. value       0.2 = laboratory reporting limit (RL)         = mean       15 = no. samples at RL         = standard deviation       RL is limit of detection of the method user         ests       • test         = t_0       -1749.76 = k_0         = t_{(n-1,0.95)}       4.36 = k_{0.05}         = 95% UCL (US ₉₅ )       1.454144 = 95% UCL (US ₉₅ )         ignificance test at 95% confidence level       is (H_0) = level of contamination is the same as, or higher than, the GAC botheses (H_1) = level of contamination is lower than the GAC treated as non-normally distributed         chever Theorem - Ho rejected, true mean <=GAC
			3 = 0.583478 = 0.957699 = <b>Statistical te</b> <b>One-sample</b> -1749.759 = 1.717 = 0.926353 = <b>Results of s</b> Null hypothesi Alternative hypothesi Alternative hypothesi Alternative hypothesi Data set Therefore: <b>Use Chebyo</b> <b>US</b> ₉₅ =	= max. value $0.2$ = laboratory reporting limit (RL)         = mean       15 = no. samples at RL         = standard deviation       RL is limit of detection of the method user         ests <b>One-sided Chebychev Theorem</b> = t ₀ -1749.76 = k ₀ = t _(n-1,0.95) 4.36 = k _{0.05} = 95% UCL (US ₉₅ )       1.454144 = 95% UCL (US ₉₅ )         ignificance test at 95% confidence level       is (H ₀ ) = level of contamination is the same as, or higher than, the GAC botheses (H ₁ ) = level of contamination is lower than the GAC treated as non-normally distributed         they Theorem - Ho rejected, true mean <=GAC
			3 = 0.583478 = 0.957699 = <b>Statistical te</b> <b>One-sample</b> -1749.759 = 1.717 = 0.926353 = <b>Results of s</b> Null hypothesi Alternative hypothesi Alternative hypothesi <b>Ste chebyc</b> <b>USe Chebyc</b> <b>US</b> ₉₅ =	= max. value $0.2 =$ laboratory reporting limit (RL)= mean15 = no. samples at RL= standard deviationRL is limit of detection of the method usedests <b>One-sided Chebychev Theorem</b> = t_0-1749.76 = k_0= t_{(n-1,0.95)}4.36 = k_{0.05}= 95% UCL (US ₉₅ )1.454144 = 95% UCL (US ₉₅ )
			3 = 0.583478 = 0.957699 = Statistical te One-sample -1749.759 = 1.717 = 0.926353 = Null hypothes Alternative hypothes Alternative hypothes Data set Therefore: Use Chebyc US ₉₅ = Site referen Data set:	= max. value $0.2 =$ laboratory reporting limit (RL)= mean15 = no. samples at RL= standard deviationRL is limit of detection of the method usedests <b>Che-sided Chebychev Theorem</b> = t_0-1749.76 = k_0= t_{(n-1,0.95)}4.36 = k_{0.05}= 95% UCL (US ₉₅ )1.454144 = 95% UCL (US ₉₅ )ignificance test at 95% confidence levelis (H_0) = level of contamination is the same as, or higher than, the GACbotheses (H_1) = level of contamination is lower than the GACtreated as non-normally distributedthey Theorem - Ho rejected, true mean <=GAC
			3 = 0.583478 = 0.957699 = <b>Statistical te</b> <b>One-sample</b> -1749.759 = 1.717 = 0.926353 = <b>Results of s</b> Null hypothesy Alternative hyp Data set <b>Therefore:</b> <b>Use Chebyo</b> <b>US</b> ₉₅ = <b>Site referen</b> Data set: I Client:	= max. value $0.2 =$ laboratory reporting limit (RL)= mean15 = no. samples at RL= standard deviationRL is limit of detection of the method usedests <b>One-sided Chebychev Theorem</b> = t_0-1749.76 = k_0= t_{(n-1,0.95)}4.36 = k_{0.05}= 95% UCL (US ₉₅ )1.454144 = 95% UCL (US ₉₅ )ignificance test at 95% confidence levelis (H_0) = level of contamination is the same as, or higher than, the GACtreated as non-normally distributedthev Theorem - Ho rejected, true mean <=GAC
			3 = 0.583478 = 0.957699 = <b>Statistical te</b> <b>One-sample</b> -1749.759 = 1.717 = 0.926353 = <b>Results of s</b> Null hypothesy Alternative hyp Data set <b>Therefore:</b> <b>Use Chebyo</b> <b>US</b> ₉₅ = <b>Site referen</b> Data set: I Client: I Site:	= max. value       0.2 = laboratory reporting limit (RL)         = mean       15 = no. samples at RL         = standard deviation       RL is limit of detection of the method used         ests       etest         = t ₀ -1749.76 = k ₀ = t ₀ 4.36 = k _{0.05} = 95% UCL (US ₉₅ )       1.454144 = 95% UCL (US ₉₅ )         ignificance test at 95% confidence level       is (H ₀ ) = level of contamination is the same as, or higher than, the GAC potheses (H ₁ ) = level of contamination is lower than the GAC treated as non-normally distributed         they Theorem - Ho rejected, true mean <=GAC
			3 = 0.583478 = 0.957699 = <b>Statistical te</b> <b>One-sample</b> -1749.759 = 1.717 = 0.926353 = <b>Results of s</b> Null hypothesy Alternative hyp Data set <b>Therefore:</b> <b>Use Chebyo</b> <b>US</b> ₉₅ = <b>Site referen</b> Data set: I Client: I Site: I Job no: 0	= max. value $0.2 =$ laboratory reporting limit (RL)= mean15 = no. samples at RL= standard deviationRL is limit of detection of the method usedestschest= to-1749.76 = k_0= t(n-1,0.95)4.36 = k_{0.05}= 95% UCL (US ₉₅ )1.454144 = 95% UCL (US ₉₅ )ignificance test at 95% confidence levelis (H ₀ ) = level of contamination is the same as, or higher than, the GACtreated as non-normally distributedthev Theorem - Ho rejected, true mean <=GAC









Chemical and data (mg/kg)	Moon Aba		ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL) (red denotes >= GAC)		<b>olute Deviation for p</b> <b>5</b> = critical value of to	
Cyanide (free)	Potential	Sample	Visual assessment - Q-g & histogram plots
0.5	Outlier? n/a	HTP15 @ 0.25	Q-q plot
0.5	n/a	HTP17 @ 0.15	
0.5	n/a	HTP20 @ 0.15	
0.5	n/a	HTP22 @ 0.20	3
0.5	n/a	HTP24 @ 0.20	
0.5	n/a	HTP25 @ 0.15	
0.5	n/a	HTP26 @ 0.10	
0.5	n/a	HTP29 @ 0.15	
0.5	n/a n/a	HTP33 @ 0.25 HTP39 @ 0.20	
0.5	n/a	HTP40 @ 0.20	
0.5	n/a	HTP41 @ 0.1	
0.5	n/a	HTP43 @ 0.2	
0.5	n/a	HTP44 @ 0.7	
0.5	n/a	HTP45 @ 0.4	
0.5	n/a	HTP46 @ 0.7	-2
5	n/a	TP5 @ 0.1	
5	n/a n/a	TP8 @ 0.4 TP8 @ 2.8	
5	n/a n/a	TP9 @ 0.6	Theoretical (zi)
	11/4	11 3 @ 0.0	
			Cyanide (free)
			-
			16
			14 +
			1212
			-10 -5 0 5 10
			<b><u> </u></b>
			2 Concentration
			Data Normal distribution
			anna
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			750 = GAC (critical conc.) (mg/kg)
			20 = no.  samples $0 = no.  samples > or = GAC$
			0.5 = min. value
			5 = max. value 0.5 = laboratory reporting limit (RL) 1.4 = mean 16 = no. samples at RL
			1.846761 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-1812.818 = t_0$ $-1812.82 = k_0$
			$1.729 = t_{(n-1,0.95)} \qquad \qquad 4.36 = k_{0.05}$
			2.113988 = 95% UCL (US ₉₅ ) 3.200455 = 95% UCL (US ₉₅ )
			Results of significance test at 95% confidence level
			Null hypothesis $(H_0)$ = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses ( $H_1$ ) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
· · · · · · · · · · · · · · · · · · ·			Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC
			$US_{95} = 3.2004547$ GAC = 750 (US95 = 0.004 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: MRB & WMF
		1	Client: Bovis Barratt and Taylor Wimpy
		1	Site: Land at Bankside, Banbury
	1		
			Job no: C12702

Chemical and data (mg/kg) (blue denotes <= RL)		STATISTICAL	
(red denotes >= GAC)	Potential		
Phenol (total)	Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.3	n/a	HTP15 @ 0.25	<u>Q-q plot</u>
0.3	n/a	HTP17 @ 0.15	
0.3	n/a	HTP20 @ 0.15	3
0.3	n/a	HTP22 @ 0.20	
0.3	n/a	HTP24 @ 0.20 HTP25 @ 0.15	
0.3	n/a n/a	HTP26 @ 0.15	2
0.3	n/a	HTP29 @ 0.15	
0.3	n/a	HTP33 @ 0.25	
0.3	n/a	HTP39 @ 0.20	
0.3	n/a	HTP40 @ 0.20	
0.01	n/a	HTP41 @ 0.1	Sample (j)
0.1	n/a	HTP43 @ 0.2	_
0.044	n/a	HTP44 @ 0.7	
0.054	n/a	HTP45 @ 0.4	
0.091	n/a	HTP46 @ 0.7	-2
0.5	n/a	TP5 @ 0.1 TP8 @ 0.4	
1.1	n/a n/a	TP8 @ 0.4 TP8 @ 2.8	
0.5	n/a	TP9 @ 0.6	Theoretical (zi)
			Phenol (total)
			_
			strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation strementation streme
			2
			- <b>6</b> -1 -0.5 0 0.5 1 1.5
			Z Concentration
			Concentration
			_
			Data      Normal distribution
			mul
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			290 = GAC (critical conc.) (mg/kg)
			20 = no.  samples $0 = no.  samples  > or = GAC$
			0.01 = min. value
			1.1 = max. value0.3 = laboratory reporting limit (RL)0.30995 = mean16 = no. samples at RL
			0.235289 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-5506.148 = t_0$ $-5506.15 = k_0$
			$1.729 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			0.400916 = 95% UCL (US ₉₅ ) 0.539339 = 95% UCL (US ₉₅ )
			Results of significance test at 95% confidence level
			Null hypothesis $(H_0)$ = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
			Data set treated as <b>non-normally distributed</b>
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			US ₉₅ = 0.5393388 GAC = 290 (US95 = 0.002 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: MRB & WMF Client: Bovis Barratt and Taylor Wimpy
			Client: Bovis Barratt and Taylor Wimpy Site: Land at Bankside, Banbury
			Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008.Guidance on comparing soil contamination with a critical concentration.
L	1		.,

(red denote	otes <= RL) es >= GAC)		blute Deviation for po = critical value of te	
	,	3.5 Potential		
Acena	phthene	Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.	.01	n/a	HTP15 @ 0.25	<u>Q-q plot</u>
0.	.01	n/a	HTP17 @ 0.15	
	.01	n/a	HTP20 @ 0.15	
	.01	n/a	HTP22 @ 0.20	
	.01	n/a	HTP24 @ 0.20	
	.01 .01	n/a n/a	HTP25 @ 0.15 HTP26 @ 0.10	
	.01	n/a	HTP28 @ 0.10	
	.01	n/a	HTP33 @ 0.25	
	.01	n/a	HTP39 @ 0.20	
	.01	n/a	HTP40 @ 0.20	
0.	.01	n/a	HTP41 @ 0.1	
С	).1	n/a	HTP43 @ 0.2	
	044	n/a	HTP44 @ 0.7	-1
	054	n/a	HTP45 @ 0.4	
	091	n/a	HTP46 @ 0.7	-2
	0.5	n/a	TP5 @ 0.1	
	).5	n/a	TP8 @ 0.4	
	).5 ).5	n/a	TP8 @ 2.8 TP9 @ 0.6	Theoretical (zi)
0	J.O	n/a	1179 @ 0.0	
				Acenaphthene
				_
				- 12 -
				stremensee -1 -0.5 0 0.5 1
				2
				<b>– E</b> -1 -0.5 0 0.5 1
				Z Concentration
				Concentration
				-
				ww
				DataNormal distribution
				•••
				Basic data Risk parameter
				Human health - residential with plant uptake (2.5%SOM)
				480 = GAC (critical conc.) (mg/kg)
				20 = no.  samples $0 = no.  samples > or = GAC$
				$0.01 = \min \text{ value}$
				0.5 = max. value 0.01 = laboratory reporting limit (RL) 0.12045 = mean 12 = no. samples at RL
				0.12045 = mean 12 = n0. samples at RL 0.196572 = standard deviation RL is limit of detection of the method used
				Statistical tests
				One-sample t-test One-sided Chebychev Theorem
				$-10917.57 = t_0$ $-10917.6 = k_0$
				$1.729 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
				$0.196448 = 95\% \text{ UCL (US}_{95}) \qquad 0.312093 = 95\% \text{ UCL (US}_{95})$
		1		Results of significance test at 95% confidence level
				Null hypothesis ( $H_0$ ) = level of contamination is the same as, or higher than, the GAC
		-		Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
				Data set treated as non-normally distributed
				Therefore:
				Use Chebychev Theorem - Ho rejected, true mean <=GAC
				US ₉₅ = 0.3120929 GAC = 480 (US95 = 0.001 x GAC)
				Site reference POTENTIALLY SUITABLE FOR USE
				Data set: MRB & WMF
				Client: Bovis Barratt and Taylor Wimpy
				Site: Land at Bankside, Banbury
				Job no: C12702
		-		Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.

	Chemical and data (mg/kg) (blue denotes <= RL) (red denotes >= GAC)		STATISTICAL blute Deviation for p = critical value of te	otential ou	tliers	ONMENTAL SOIL DATA able as 50% or more of values are the same.
		Potential				
	Acenaphthylene	Outlier?	Sample	Visua	assessment - Q-q & histor	
	0.01	n/a	HTP15 @ 0.25		<u>Q</u> .	<u>-q plot</u>
	0.01	n/a	HTP17 @ 0.15			
	0.01	n/a	HTP20 @ 0.15			2
	0.01	n/a	HTP22 @ 0.20			3
	0.01	n/a	HTP24 @ 0.20			
	0.01	n/a n/a	HTP25 @ 0.15 HTP26 @ 0.10			2
	0.01	n/a	HTP29 @ 0.10			
	0.01	n/a	HTP33 @ 0.25	(c-		-1
	0.073	n/a	HTP39 @ 0.20	Sample (yi)		
~~~~~	0.026	n/a	HTP40 @ 0.20	<u>e</u>		
	0.01	n/a	HTP41 @ 0.1	Ē.	3 -2 1	
	0.063	n/a	HTP43 @ 0.2	Sa		
	0.03	n/a	HTP44 @ 0.7			-1
	0.028	n/a	HTP45 @ 0.4			
	0.01	n/a	HTP46 @ 0.7			-2
	0.5	n/a	TP5 @ 0.1			
	0.5	n/a	TP8 @ 0.4			-3
	0.5	n/a	TP8 @ 2.8		Th	eoretical (zi)
	0.5	n/a	TP9 @ 0.6		110	
					<u>Acer</u>	haphthylene
						16
						14
				s		12
				measurements		10
				Ĕ		
				2		8
				eas		6
						4
				Number of		
						2
				<u>E</u>		
				ź	-1 -0.5	0 0.5 1
					c	Concentration
					Dete	
					Data	Normal distribution
				<u> </u>		
				Basic d		parameter
						al with plant uptake (2.5%SOM)
					00 = GAC (critical conc.) (n	
				^	20 = no. samples .01 = min. value	0 = no. samples > or = GAC
					0.5 = max. value	0.01 = laboratory reporting limit (RL)
					65 = mean	11 = no. samples at RL
					526 = standard deviation	RL is limit of detection of the method used
				Statistic	al tests	
				One-sa	nple t-test	One-sided Chebychev Theorem
				-9053.6	$566 = t_0$	$-9053.67 = k_0$
				1.7	$729 = t_{(n-1,0.95)}$	$4.36 = k_{0.05}$
					867 = 95% UCL (US ₉₅)	0.309073 = 95% UCL (US ₉₅)
			1			
					of significance test at 95%	
						ion is the same as, or higher than, the GAC
						tamination is lower than the GAC
					a set treated as non-normall	y distributed
				Therefor		
					ebychev Theorem - Ho reject	
					₅ = 0.309073 GAC = 400	(US95 = 0.001 x GAC)
1				Site ref		POTENTIALLY SUITABLE FOR USE
				Data set		• / •
				Client	Bovis Barratt and Taylor V	(Imp)/
				Client:		
				Site:	Land at Bankside, Banbu	
					Land at Bankside, Banbu C12702	

(mg/kg) (blue denotes <= RL)

(red denotes >= GAC)

Anthracene

0.01 0.01

0.11

0.01

0.01

0.01

0.01

0.012

0.01

0.14 0.064

0.03

0.02

0.01

0.01

0.031 0.5

0.78

0.5

0.5

Potential

Outlier?

Yes

Yes

Yes

Yes

Yes

Yes

Yes

Sample

HTP15 @ 0.25

HTP17 @ 0.15 HTP20 @ 0.15

HTP22 @ 0.20

HTP24 @ 0.20

HTP25 @ 0.15

HTP26 @ 0.10

HTP29 @ 0.15 HTP33 @ 0.25

HTP39 @ 0.20

HTP40 @ 0.20

HTP41 @ 0.1

HTP43 @ 0.2

HTP44 @ 0.7 HTP45 @ 0.4

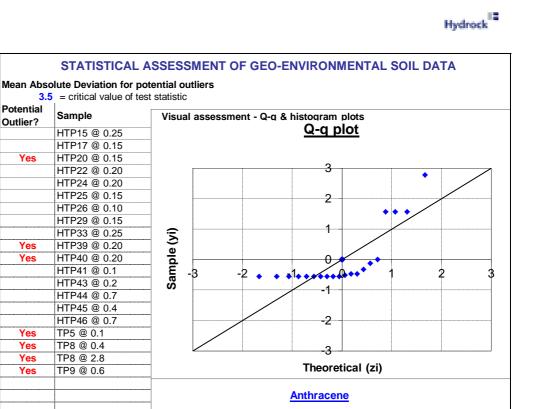
HTP46 @ 0.7

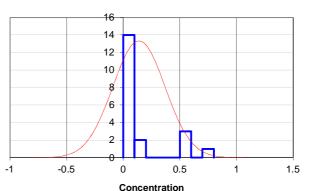
TP5 @ 0.1

TP8 @ 0.4

TP8 @ 2.8

TP9 @ 0.6





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		ž -1 -0.5 (0 0.5 1 1.5
			Concentration
		Data	Normal distribution
			parameter
		Human health - residentia	al with plant uptake (2.5%SOM)
		4900 = GAC (critical conc.) (n	ng/kg)
		20 = no. samples	0 = no. samples > or = GAC
		0.01 = min. value	
		0.78 = max. value	0.01 = laboratory reporting limit (RL)
		0.13885 = mean	9 = no. samples at RL
		0.230778 = standard deviation	RL is limit of detection of the method used
		Statistical tests	
		One-sample t-test	One-sided Chebychev Theorem
		$-94951.9 = t_0$	$-94951.9 = k_0$
		$1.729 = t_{(n-1,0.95)}$	$4.36 = k_{0.05}$
		0.228073 = 95% UCL (US ₉₅)	0.363842 = 95% UCL (US ₉₅)
1		•	•
		Results of significance test at 95%	confidence level
			ion is the same as, or higher than, the GAC
		Alternative hypotheses (H ₁) = level of con	tamination is lower than the GAC
		Data set treated as non-normal	y distributed
		Therefore:	
 	· · ·	Lies Chebyshey Theorem . He reis	ated true mean -CAC

С Use Chebychev Theorem - Ho rejected, true mean <=GAC US₉₅ = 0.3638418 GAC = 4900 (US95 = 0 x 0 $(US95 = 0 \times GAC)$ Site reference SUIT Data set: MRB & WMF Client: Bovis Barratt and Taylor Wimpy Site[.] Land at Bankside, Banbury Job no: C12702 Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.

Chemical and data (mg/kg)			ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(blue denotes <= RL) (red denotes >= GAC)		olute Deviation for p 5 = critical value of to	
	Potential		
Benz(a)anthracene	Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.01		HTP15 @ 0.25	<u>Q-q plot</u>
0.01		HTP17 @ 0.15	
0.36	Yes	HTP20 @ 0.15	
0.01		HTP22 @ 0.20	
0.01		HTP24 @ 0.20	
0.01		HTP25 @ 0.15	
0.047		HTP26 @ 0.10	
0.099		HTP29 @ 0.15	
0.023		HTP33 @ 0.25	Beneficial Structure (iv) and the second structure (iv) and the se
0.95	Yes	HTP39 @ 0.20	
0.48	Yes	HTP40 @ 0.20	
0.11		HTP41 @ 0.1	
0.04		HTP43 @ 0.2	0
0.01		HTP44 @ 0.7 HTP45 @ 0.4	
0.01		HTP46 @ 0.7	
0.5	Yes	TP5 @ 0.1	
0.5	Yes	TP8 @ 0.4	
0.5	Yes	TP8 @ 2.8	
0.5	Yes	TP9 @ 0.6	Theoretical (zi)
0.0		1	
			Benz(a)anthracene
			strengen v strengen v streng
			- - - - - - - - - -
			_
			Concentration
			••••
			Data Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			4.7 = GAC (critical conc.) (mg/kg)
			20 = no. samples $0 = no. samples > or = GAC$
			0.01 = min. value
			0.01 = laboratory reporting limit (RL)
			0.22245 = mean 8 = no. samples at RL
			0.29267 = standard deviation RL is limit of detection of the method use
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-68.41908 = t_0$ $-68.4191 = k_0$
			$1.729 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			$\begin{array}{c} -1.125 = -(n_{1,0.95}) \\ 0.335601 = 95\% \text{ UCL } (\text{US}_{95}) \\ 0.507781 = 95\% \text{ UCL } (\text{US}_{95}) \end{array}$
			0.00001 - 30% 001 (0095) = 30% 001 (0095)
			Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses (H_1) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			$US_{95} = 0.5077815$ GAC = 4.7 (US95 = 0.108 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
			Data set: MRB & WMF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration

Chemical and data (mg/kg) (blue denotes <= RL)	Moon Ab-	STATISTICAL plute Deviation for p	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
(red denotes >= GAC)	3.5	\mathbf{b}_{i} = critical value of te	
Benzo(a)pyrene	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.01		HTP15 @ 0.25	<u>Q-q plot</u>
0.01		HTP17 @ 0.15	
0.38	Yes	HTP20 @ 0.15	
0.01		HTP22 @ 0.20	3
0.01		HTP24 @ 0.20	
0.01		HTP25 @ 0.15	2
0.01		HTP26 @ 0.10	
0.07		HTP29 @ 0.15	
0.018		HTP33 @ 0.25	
0.69	Yes	HTP39 @ 0.20	
0.33	Yes	HTP40 @ 0.20	
0.055		HTP41 @ 0.1	Eg3 -2 -1 _0 → 1 2 3
0.023		HTP43 @ 0.2	/Ő
0.01		HTP44 @ 0.7	
0.01		HTP45 @ 0.4	
0.01		HTP46 @ 0.7	
0.5	Yes	TP5 @ 0.1	
0.95	Yes	TP8 @ 0.4	
0.5	Yes	TP8 @ 2.8	Theoretical (zi)
0.5	Yes	TP9 @ 0.6	
			Benzo(a)pyrene
			12
			- $ -$
			Ž -1.5 -1 -0.5 0 0.5 1 1.5
			Concentration
			Data Normal distribution
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			0.94 = GAC (critical conc.) (mg/kg)
			20 = no. samples 1 = no. samples > or = GAC
			0.01 = min. value
			0.95 = max. value 0.01 = laboratory reporting limit (RL)
			0.2053 = mean 9 = no. samples at RL
			0.285493 = standard deviation RL is limit of detection of the method used
			Statistical tasta
			Statistical tests
			One-sided Chebychev Theorem
			$-11.5088 = t_0$ $-11.5088 = k_0$
			$1.729 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			$0.315676 = 95\% \text{ UCL } (\text{US}_{95})$ $0.483634 = 95\% \text{ UCL } (\text{US}_{95})$
			Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses (H ₁) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			US ₉₅ = 0.4836342 GAC = 0.94 (US95 = 0.515 x GAC)
	-		Site reference POTENTIALLY SUITABLE FOR USE
			Data set: MRB & WMF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			Job no: C12702
			Reference: CL:AIRE & CIEH. May 2008.Guidance on comparing soil contamination with a critical concentration.
<u>,</u>		•	

(mg/kg)

(blue denotes <= RL) (red denotes >= GAC)

Benzo(b)fluoranthene

0.01

0.01 0.5

0.01

0.01 0.01

0.046 0.12

0.021

1.2

0.62

0.18

0.056

0.01

0.01

0.01

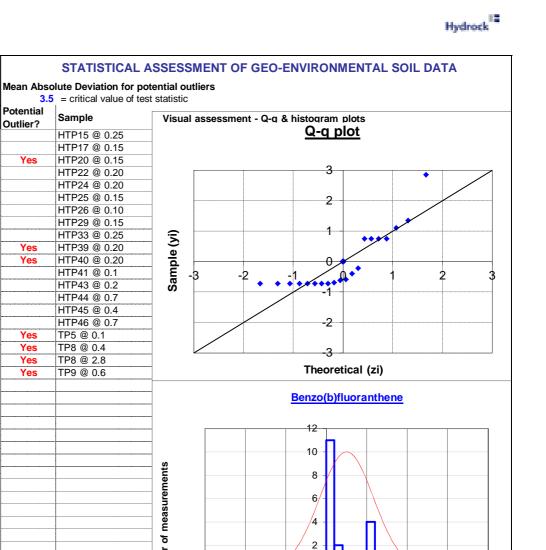
Potential

Outlier?

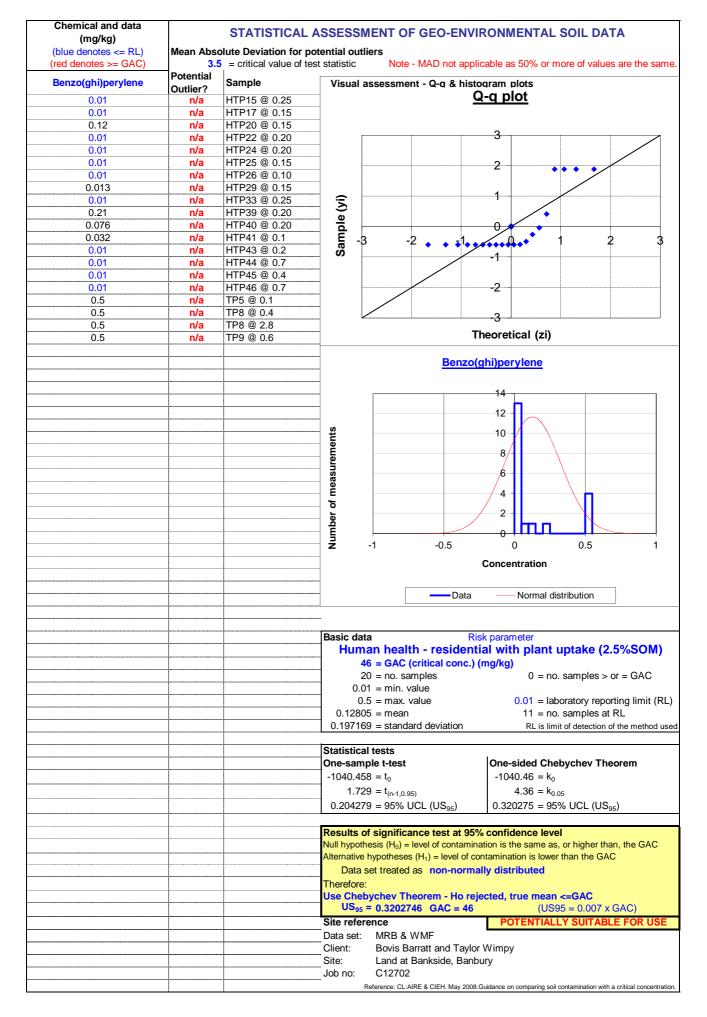
Yes

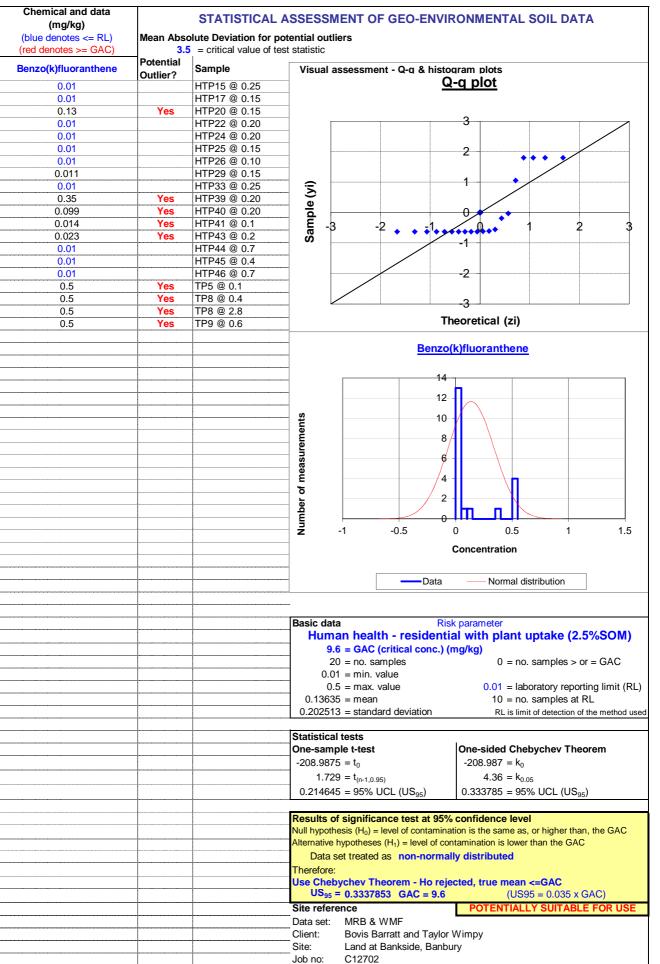
Yes

Yes



0.01		HTP46 @ 0.7	-2
0.5	Yes	TP5 @ 0.1	
0.7	Yes	TP8 @ 0.4	
0.5	Yes	TP8 @ 2.8	
0.5	Yes	TP9 @ 0.6	Theoretical (zi)
			Benzo(b)fluoranthene
			<u>v</u>
			stremen streme
			Concentration
			Data Normal distribution
			new Control of Control
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			6.5 = GAC (critical conc.) (mg/kg)
			20 = no. samples $0 = no. samples > or = GAC$
			0.01 = min. value
			1.2 = max. value $0.01 = laboratory reporting limit (RL)$
			0.25115 = mean 8 = no. samples at RL
			0.332943 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-83.93547 = t_0$ $-83.9355 = k_0$
			$1.729 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			0.379871 = 95% UCL (US ₉₅) $0.575744 = 95%$ UCL (US ₉₅)
			Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses (H_1) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
		+	Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			$US_{95} = 0.5757444$ GAC = 6.5 (US95 = 0.089 x GAC)
			Site reference POTENTIALLY SUITABLE FOR USE
		+	Data set: MRB & WMF
		+	Client: Bovis Barratt and Taylor Wimpy
		+	Site: Land at Bankside, Banbury
			Job no: C12702
			COD TIO: CT2TO2 Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.



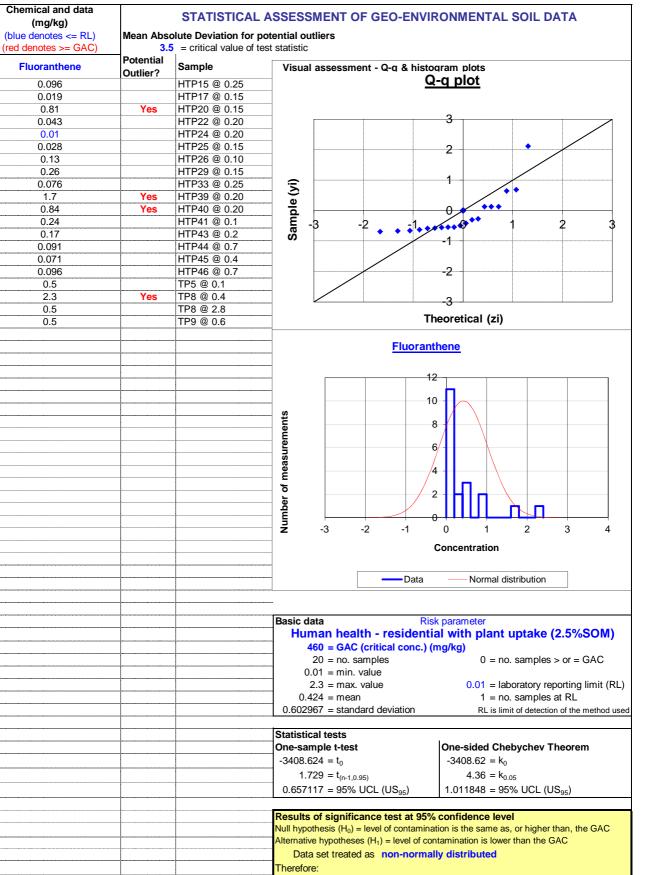


Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.



	Chemical and data (mg/kg) (blue denotes <= RL)	Mean Abso	STATISTICAL	ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA
	(red denotes >= GAC)	3.5	= critical value of t	
	Chrysene	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots
	0.01	outlier?	HTP15 @ 0.25	<u>Q-q plot</u>
	0.01		HTP17 @ 0.15	
	0.3	Yes	HTP20 @ 0.15	<u>^</u>
	0.01		HTP22 @ 0.20	
	0.01		HTP24 @ 0.20	
	0.01 0.029		HTP25 @ 0.15 HTP26 @ 0.10	
	0.025		HTP29 @ 0.15	
	0.024		HTP33 @ 0.25	
	0.78	Yes	HTP39 @ 0.20	
	0.32	Yes	HTP40 @ 0.20	
	0.11 0.054		HTP41 @ 0.1	$ \frac{1}{1}$ $\frac{1}{2}$ $\frac{1}{3}$
	0.054		HTP43 @ 0.2 HTP44 @ 0.7	-
	0.01		HTP45 @ 0.4	
	0.01		HTP46 @ 0.7	-2
	0.5	Yes	TP5 @ 0.1	
	0.72	Yes	TP8 @ 0.4	-3
	0.5	Yes	TP8 @ 2.8	Theoretical (zi)
	0.5	Yes	TP9 @ 0.6	
				Chrysene
				_
				- \$ 10 -
				Base Contraction of the second
				0.5 0 0.5 1 1.5
				Concentration
				Concentration
				Data Normal distribution
				Basic data Risk parameter
		-		Human health - residential with plant uptake (2.5%SOM)
		-		8 = GAC (critical conc.) (mg/kg)
				20 = no. samples 0 = no. samples > or = GAC
				0.01 = min. value
				0.78 = max. value 0.01 = laboratory reporting limit (RI
				0.19985 = mean 8 = no. samples at RL 0.260973 = standard deviation RL is limit of detection of the method us
				Statistical tests
				One-sample t-test One-sided Chebychev Theorem
				$-133.6664 = t_0$ $-133.666 = k_0$
				$1.729 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
				$0.300746 = 95\% \text{ UCL (US}_{95})$ $0.454279 = 95\% \text{ UCL (US}_{95})$
				Results of significance test at 95% confidence level
				Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H_1) = level of contamination is lower than the GAC
				Data set treated as non-normally distributed
				Therefore:
				Use Chebychev Theorem - Ho rejected, true mean <=GAC
				US ₉₅ = 0.4542794 GAC = 8 (US95 = 0.057 x GAC)
				Site reference POTENTIALLY SUITABLE FOR USE
				Data set: MRB & WMF
				Client: Bovis Barratt and Taylor Wimpy
1 7				Site: Land at Bankside, Banbury
				Job no: C12702 Reference: CL:AIRE & CIEH. May 2008.Guidance on comparing soil contamination with a critical concentrat

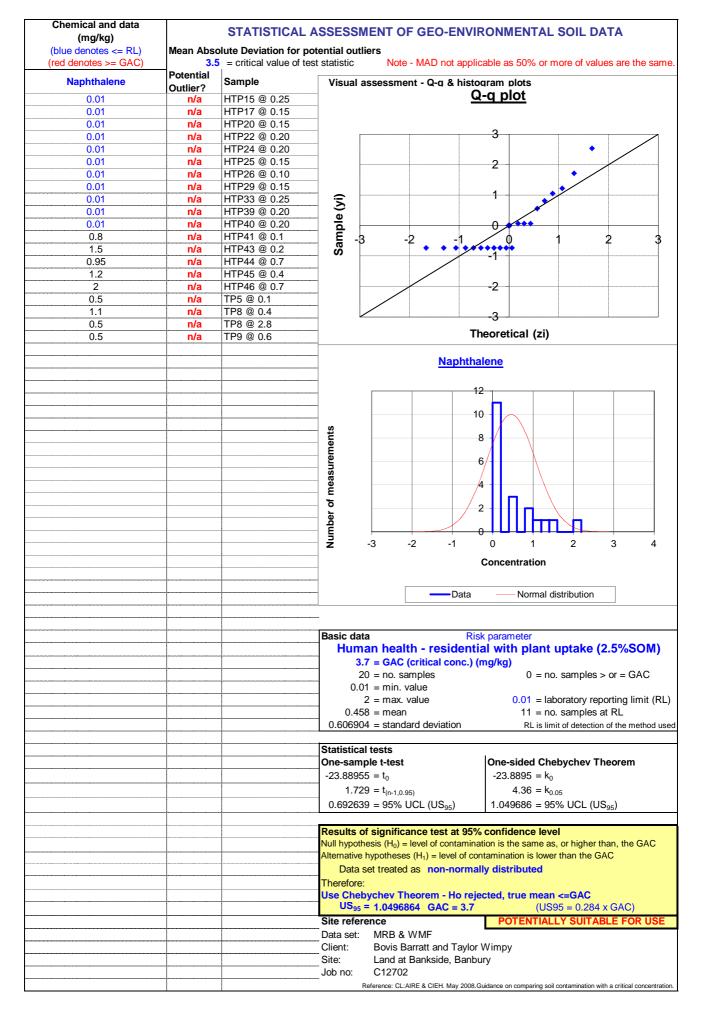
Chemical and data (mg/kg) (blue denotes <= RL)		olute Deviation for p	
(red denotes >= GAC)		5 = critical value of te	st statistic Note - MAD not applicable as 50% or more of values are the same.
Dibenz(a,h)anthracene	Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.01	n/a	HTP15 @ 0.25	<u>Q-q plot</u>
0.01	n/a	HTP17 @ 0.15	
0.01	n/a	HTP20 @ 0.15	
0.01	n/a	HTP22 @ 0.20	3
0.01	n/a	HTP24 @ 0.20	
0.01	n/a	HTP25 @ 0.15	
0.01	n/a	HTP26 @ 0.10	
0.01	n/a n/a	HTP29 @ 0.15 HTP33 @ 0.25	
0.01	n/a	HTP39 @ 0.20	- (j)
0.01	n/a	HTP40 @ 0.20	
0.01	n/a	HTP41 @ 0.1	
0.01	n/a	HTP43 @ 0.2	
0.01	n/a	HTP44 @ 0.7	-1
0.01	n/a	HTP45 @ 0.4	
0.01	n/a	HTP46 @ 0.7	-2
0.5	n/a	TP5 @ 0.1	
0.5	n/a	TP8 @ 0.4	
0.5	n/a n/a	TP8 @ 2.8 TP9 @ 0.6	Theoretical (zi)
0.5	TI/a		
			Dibenz(a,h)anthracene
			16
			14 -
			12/ 10- 8- 6-
			_ ზ 4
			Ž −1 −0.5 0 0.5 1
			Concentration
			Data Normal distribution
			ww
			Basic data Risk parameter
			Human health - residential with plant uptake (2.5%SOM)
			0.86 = GAC (critical conc.) (mg/kg)
	-		20 = no. samples $0 = no. samples > or = GAC$
	-		0.01 = min. value
			0.5 = max. value 0.01 = laboratory reporting limit (RL)
			0.108 = mean 16 = no. samples at RL
			0.201092 = standard deviation RL is limit of detection of the method used
			Statistical tests
			One-sample t-test One-sided Chebychev Theorem
			$-16.72394 = t_0$ $-16.7239 = k_0$
			$1.729 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$
			0.185745 = 95% UCL (US ₉₅) $0.30405 = 95%$ UCL (US ₉₅)
			Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC Alternative hypotheses (H_1) = level of contamination is lower than the GAC
		-	Data set treated as non-normally distributed
	-		Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
	1		$US_{95} = 0.3040495$ GAC = 0.86 (US95 = 0.354 x GAC)
	-		Site reference POTENTIALLY SUITABLE FOR USE
		1	Data set: MRB & WMF
			Client: Bovis Barratt and Taylor Wimpy
			Site: Land at Bankside, Banbury
			Job no: C12702
	1	1	Reference: CL:AIRE & CIEH. May 2008.Guidance on comparing soil contamination with a critical concentration.

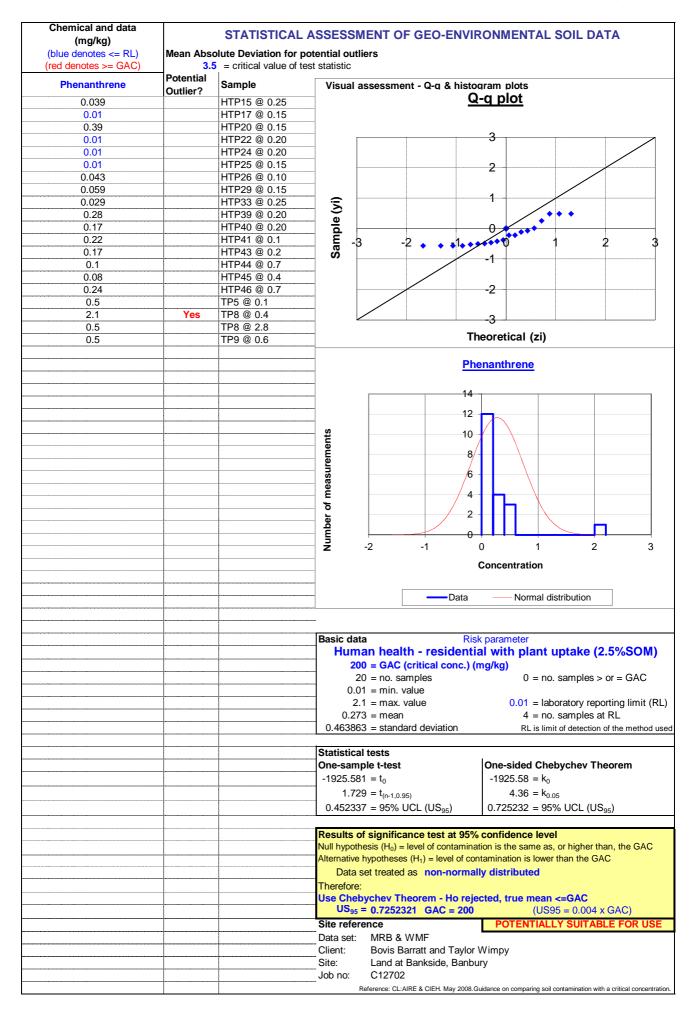




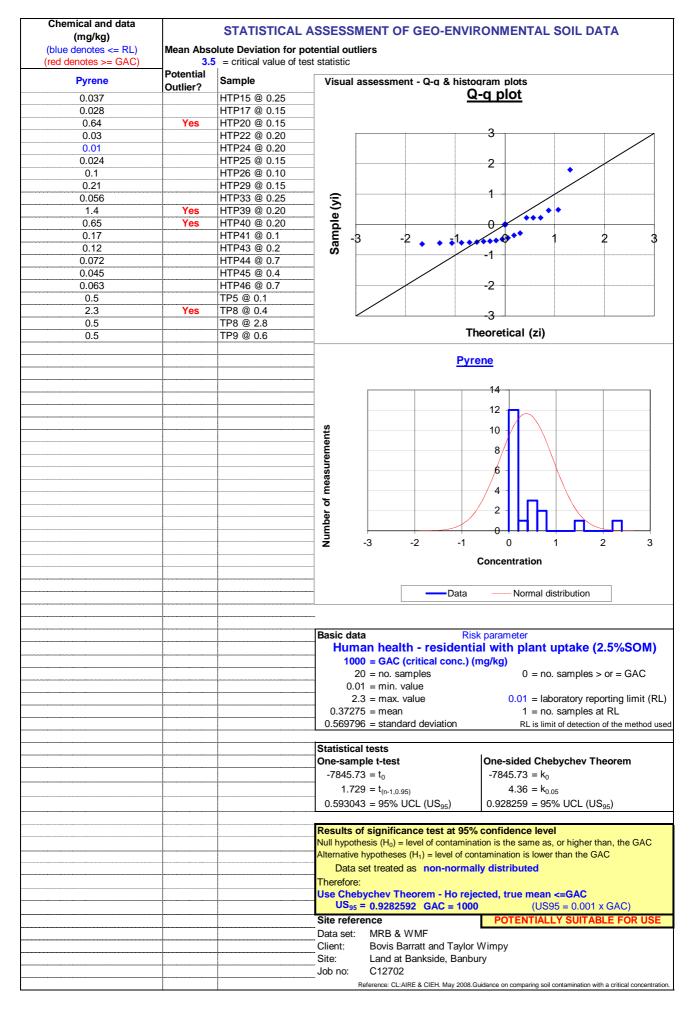
Chemical and data (mg/kg) (blue denotes <= RL)	STATISTICAL ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA Mean Absolute Deviation for potential outliers					
(red denotes >= GAC)	3.5 Potential	= critical value of to	est statistic	Note - MAD not applica	able as 50% or more of values are the same.	
Fluorene	Outlier?	Sample	Visual assessment - Q-q & histogram plots			
0.01	n/a	HTP15 @ 0.25		<u>Q</u> .	<u>-q plot</u>	
0.01	n/a	HTP17 @ 0.15				
0.01	n/a	HTP20 @ 0.15			-3	
0.01	n/a	HTP22 @ 0.20 HTP24 @ 0.20			3	
0.01	n/a n/a	HTP24 @ 0.20				
0.01	n/a	HTP26 @ 0.10			2	
0.01	n/a	HTP29 @ 0.15				
0.01	n/a	HTP33 @ 0.25	ii			
0.01	n/a	HTP39 @ 0.20	Sample (yi)			
0.01	n/a n/a	HTP40 @ 0.20 HTP41 @ 0.1	Id			
0.08	n/a	HTP43 @ 0.2	an San	β −2 • • •1•••	••••••••••• 1 2 3	
0.038	n/a	HTP44 @ 0.7			-1	
0.053	n/a	HTP45 @ 0.4				
0.11	n/a	HTP46 @ 0.7			-2	
0.5	n/a	TP5 @ 0.1				
0.5	n/a	TP8 @ 0.4			-3	
0.5	n/a n/a	TP8 @ 2.8 TP9 @ 0.6		Th	eoretical (zi)	
0.5	n/a	1P9 @ 0.6				
				Fluo	orene	
					-14	
					12 -	
			v		10	
			measurements			
			em		8/-	
			s		6	
			lea			
					4	
			Number of		2	
			hbe			
				-1 -0.5	0 0.5 1	
			č			
				(Concentration	
				Data	Normal distribution	
			<u> </u>	<u> </u>		
			Basic da		a parameter Il with plant uptake (2.5%SOM)	
				80 = GAC (critical conc.) (n		
				20 = no. samples	0 = no. samples > or = GAC	
				01 = min. value		
			C	0.5 = max. value	0.01 = laboratory reporting limit (RL)	
				05 = mean	12 = no. samples at RL	
			0.1967	78 = standard deviation	RL is limit of detection of the method used	
			Statiati-	al tosts		
			Statistic	al tests iple t-test	One-sided Chebychev Theorem	
			-8633.4	•	$-8633.44 = k_0$	
				$42 = t_0$ 29 = t _(n-1,0.95)	$4.36 = k_{0.05}$	
		-		29 – (_{n-1,0.95)} 28 = 95% UCL (US ₉₅)	$4.30 - \kappa_{0.05}$ 0.311894 = 95% UCL (US ₉₅)	
		+	0.1301		3.571004 - 3578 OOL (0095)	
		-	Results	of significance test at 95%	confidence level	
				-	ion is the same as, or higher than, the GAC	
				Alternative hypotheses (H_1) = level of contamination is lower than the GAC		
				Data set treated as non-normally distributed		
			Therefore			
				bychev Theorem - Ho reject		
				= 0.3118943 GAC = 380	$(US95 = 0.001 \times GAC)$	
			Site refe		POTENTIALLY SUITABLE FOR USE	
			Data set:		Nimpy	
		1		Client: Bovis Barratt and Taylor Wimpy Site: Land at Bankside, Banbury		
			Job no:	C12702	,	
					idance on comparing soil contamination with a critical concentration.	
L		1		NOICICINE, OLAIRE & CIER, May 2008.GU	incense on comparing son contamination with a critical concentration	

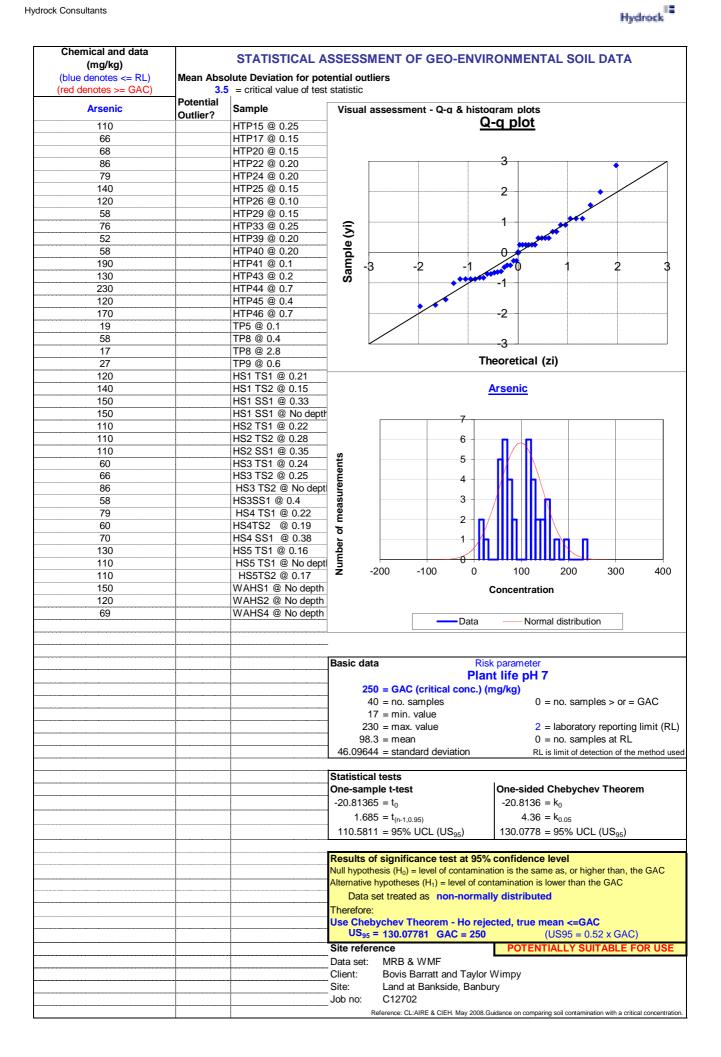
Chemical and data (mg/kg) (blue denotes <= RL)		olute Deviation for p	
(red denotes >= GAC)	3.t Potential	= critical value of te	est statistic Note - MAD not applicable as 50% or more of values are the same.
Indeno(1,2,3,cd)pyrene	Outlier?	Sample	Visual assessment - Q-q & histogram plots
0.01	n/a	HTP15 @ 0.25	<u>Q-q plot</u>
0.01	n/a	HTP17 @ 0.15	
0.083	n/a	HTP20 @ 0.15	
0.01	n/a	HTP22 @ 0.20	3
0.01	n/a	HTP24 @ 0.20	
0.01	n/a	HTP25 @ 0.15	
0.01	n/a	HTP26 @ 0.10	
0.01	n/a	HTP29 @ 0.15	
0.01	n/a	HTP33 @ 0.25	
0.22	n/a	HTP39 @ 0.20	
0.069	n/a	HTP40 @ 0.20	
0.025	n/a	HTP41 @ 0.1	
0.01	n/a	HTP43 @ 0.2	
0.01	n/a	HTP44 @ 0.7	
0.01	n/a	HTP45 @ 0.4	
0.01	n/a	HTP46 @ 0.7	-2
0.5	n/a	TP5 @ 0.1	
0.5	n/a	TP8 @ 0.4	
0.5	n/a	TP8 @ 2.8	, i i i i i i i i i i i i i i i i i i i
0.5	n/a	TP9 @ 0.6	Theoretical (zi)
			Indeno(1,2,3,cd)pyrene
			12
			- \$ 10 - 10 - 10 - 10 - 10 - 10 - 10 - 1
			┤Ē └── ↓
			ž -1 -0.5 0 0.5 1
			Concentration
			Data — Normal distribution
			Basic data Risk parameter Human health - residential with plant uptake (2.5%SOM)
			3.9 = GAC (critical conc.) (mg/kg)
			20 = no. samples $0 = no. samples > or = GAC$
			0.01 = min. value 0.5 = max. value 0.01 = laboratory reporting limit (RL)
			0.5 = max. value 0.01 = laboratory reporting limit (RL) 0.12585 = mean 12 = no. samples at RL
			0.12803 = mean 12 = n0. samples at RL 0.198031 = standard deviation RL is limit of detection of the method used
			C. 130031 - Stanuaru uzvialiuri RL IS limit of detection of the method USec
			Statistical tests
			One-sided Chebychev Theorem
			$-85.23159 = t_0$ $-85.2316 = k_0$
			$1.729 = t_{(n-1,0.95)} $
			$0.202412 = 95\% \text{ UCL } (\text{US}_{95})$ $0.318916 = 95\% \text{ UCL } (\text{US}_{95})$
			Results of significance test at 95% confidence level
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC
			Alternative hypotheses (H_1) = level of contamination is lower than the GAC
			Data set treated as non-normally distributed
			Therefore:
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			Use Chebychev Theorem - Ho rejected, true mean <=GAC US ₉₅ = 0.3189157 GAC = 3.9 (US95 = 0.082 x GAC)
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			Use Chebychev Theorem - Ho rejected, true mean <=GAC US ₉₅ = 0.3189157 GAC = 3.9 (US95 = 0.082 x GAC)
			Use Chebychev Theorem - Ho rejected, true mean <= GAC US ₉₅ = 0.3189157 GAC = 3.9 (US95 = 0.082 × GAC) Site reference POTENTIALLY SUITABLE FOR USE
			Use Chebychev Theorem - Ho rejected, true mean <=GAC
			Use Chebychev Theorem - Ho rejected, true mean <=GAC



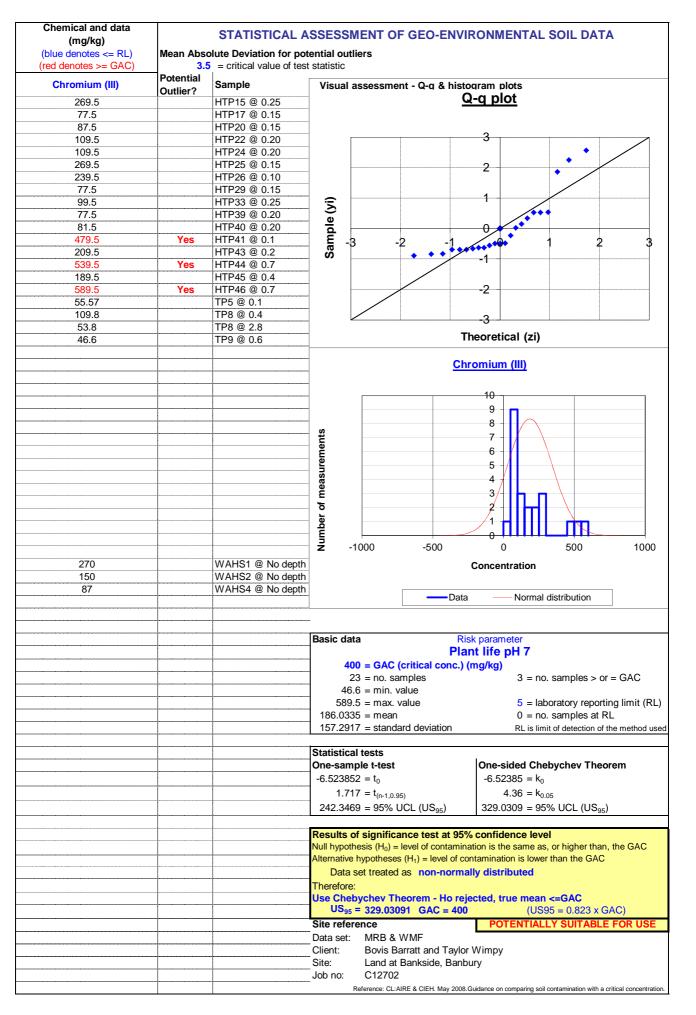








Chemical and data (mg/kg)	STATISTICAL ASSESSMENT OF GEO-ENVIRONMENTAL SOIL DATA											
(blue denotes <= RL) (red denotes >= GAC)	3.5	Mean Absolute Deviation for potential outliers 3.5 = critical value of test statistic										
Boron	Potential Outlier?	Sample	Visual assessment - Q-q & histogram plots									
0.8	Outlier	HTP15 @ 0.25	<u>Q-q plot</u>									
0.9		HTP17 @ 0.15	-									
1.4		HTP20 @ 0.15										
0.7		HTP22 @ 0.20	3									
0.9		HTP24 @ 0.20										
0.5		HTP25 @ 0.15 HTP26 @ 0.10	2									
1.3		HTP29 @ 0.15										
1.2		HTP33 @ 0.25										
0.7		HTP39 @ 0.20	Sample (i) 									
0.9		HTP40 @ 0.20										
1		HTP41 @ 0.1										
1 0.7		HTP43 @ 0.2 HTP44 @ 0.7	ο ο <u>· · · ·</u> · · · · · · · · · · · · · · ·									
1		HTP45 @ 0.4										
1.2		HTP46 @ 0.7	-2									
1.8		TP5 @ 0.1										
1		TP8 @ 0.4										
2.2	Yes	TP8 @ 2.8	Theoretical (zi)									
0.7		TP9 @ 0.6										
			Boron									
			5									
			d a a a a a a a a a a a a a									
			z -1 0 1 2 3									
			Concentration									
			Data Normal distribution									
			Basic data Risk parameter									
			Plant life pH 7									
			3 = GAC (critical conc.) (mg/kg) 20 = no. samples 0 = no. samples > or = GAC									
	-		0.5 = min. value									
	1		2.2 = max. value 0.4 = laboratory reporting limit (RL)									
			1.05 = mean 0 = no. samples at RL									
			0.400657 = standard deviation RL is limit of detection of the method used									
			Statistical tests									
			One-sided Chebychev Theorem									
			$-21.76589 = t_0$ $-21.7659 = k_0$									
			$1.729 = t_{(n-1,0.95)} \qquad 4.36 = k_{0.05}$									
			$1.204901 = 95\% \text{ UCL } (\text{US}_{95}) \qquad 1.440611 = 95\% \text{ UCL } (\text{US}_{95})$									
			Results of significance test at 95% confidence level									
			Null hypothesis (H_0) = level of contamination is the same as, or higher than, the GAC									
			Alternative hypotheses $(H_1) =$ level of contamination is lower than the GAC									
			Data set treated as non-normally distributed									
			Therefore: Use Chebychev Theorem - Ho rejected, true mean <=GAC									
			$US_{95} = 1.4406111$ GAC = 3 (US95 = 0.48 x GAC)									
			Site reference POTENTIALLY SUITABLE FOR USE									
			Data set: MRB & WMF									
			Client: Bovis Barratt and Taylor Wimpy									
			Site: Land at Bankside, Banbury									
			_Job no: C12702									
			Reference: CL:AIRE & CIEH. May 2008. Guidance on comparing soil contamination with a critical concentration.									





	Chemical and data (mg/kg) (blue denotes <= RL) (red denotes >= GAC)		STATISTICAL blute Deviation for p = critical value of te	
		Potential		
	Chromium (VI)	Outlier?	Sample	Visual assessment - Q-q & histogram plots
	0.5	n/a	HTP15 @ 0.25	<u>Q-q plot</u>
	0.5	n/a	HTP17 @ 0.15 HTP20 @ 0.15	
	0.5	n/a n/a	HTP20 @ 0.15	3
	0.5	n/a	HTP24 @ 0.20	
	0.5	n/a	HTP25 @ 0.15	2
	0.5	n/a	HTP26 @ 0.10	
	0.5	n/a	HTP29 @ 0.15	
	0.5	n/a n/a	HTP33 @ 0.25 HTP39 @ 0.20	Sample (y)
	0.5	n/a	HTP40 @ 0.20	
	0.5	n/a	HTP41 @ 0.1	
	0.5	n/a	HTP43 @ 0.2	
	0.5	n/a	HTP44 @ 0.7	
	0.5	n/a	HTP45 @ 0.4	
	0.5 0.43	n/a n/a	HTP46 @ 0.7 TP5 @ 0.1	-2
	0.2	n/a	TP8 @ 0.4	
	1.2	n/a	TP8 @ 2.8	
	0.4	n/a	TP9 @ 0.6	Theoretical (zi)
				<u>Chromium (VI)</u>
				14
				<u>\$</u> 12
				ž -0.5 0 0.5 1 1.5
				Concentration
~~~~~				Data Normal distribution
				Basic data Risk parameter
				Plant life pH 7
				25 = GAC (critical conc.) (mg/kg)
				20 = no.  samples $0 = no.  samples > or = GAC$
				0.2 = min. value 1.2 = max. value 0.5 = laboratory reporting limit (RL)
				0.5115 = mean $19 = no. samples at RL$
				0.176554 = standard deviation RL is limit of detection of the method used
				Statistical tests
				One-sample t-test         One-sided Chebychev Theorem           -620.2969 = t ₀ -620.297 = k ₀
				$\begin{array}{c} 1.729 = t_{(n-1,0.95)} \\ 0.570759 = 0.5\% (100 (115)) \\ 0.683627 = 0.5\% (101 (115)) \\ \end{array}$
			+	0.579759 = 95% UCL (US ₉₅ ) $0.683627 = 95%$ UCL (US ₉₅ )
			-	Results of significance test at 95% confidence level
				Null hypothesis ( $H_0$ ) = level of contamination is the same as, or higher than, the GAC
			-	Alternative hypotheses $(H_1)$ = level of contamination is lower than the GAC
				Data set treated as non-normally distributed
				Therefore:
				Use Chebychev Theorem - Ho rejected, true mean <=GAC
				US ₉₅ = 0.683627 GAC = 25 (US95 = 0.027 x GAC)
				Site reference POTENTIALLY SUITABLE FOR USE
				Data set: MRB & WMF Client: Bovis Barratt and Taylor Wimpy
			+	Site: Land at Bankside, Banbury
			1	
				Job no: C12702

# Summary of Remedial Targets Methodology

RTM Level 1 - Soil Zone A	ssessment	t - leachate	samples							
Water body receptor(s										
Secondary receptor(s										
	t: Tip Area	, cyclom								
	t: Bovis, Barra	att and Taylor	Wimpey Hor	nes						
	: Land at Bar	•		lico						
	: C12702	indiae, Baliba	, y							
000 110									No. Comulas	
		•	ary of Sample	<b>D</b> (		Value Being	Water Quality Target		No. Samples	
Chemicals of Potential		Summa	e Data		Compared to	(Exceede		Exceeding Wate	r	
Concern					Target =	Text)		Quality Target	Notes	
(concentrations in µg/l)	No. of	Limit of	Minimum	Maximum	95-%ile	Maximum	Inland		Inland	
						Value	Waters		Waters	
	Samples	Detection	Value	Value	Value		EQS		EQS	
Hardness as mg/l CaCO3	-	-	61	-	-	-	-			Used with some EQS.
Ag (dissolved)	2	0.5	0.5	0.5	0.5	0.5	0.05		2	EQS > LoD.
As (dissolved)	2	1	1	1.6	1.57	1.6	50		0	
B (dissolved)	2	20	20	20	20	20	2000		0	
Cd (dissolved)	2	0.08	0.08	0.08	0.08	0.08	0.09		0	
Co (dissolved)	2	1	1	1	1	1	3		0	
Cr (VI) (dissolved)	2	1	1	1	1	1	3.4		0	
Cr (III) (dissolved)	2	1	1	1	1	1	4.7		0	
Cu (dissolved)	2	1	1	6.3	6.035	6.3	6		1	
Fe (dissolved)	2	20	100	590	565.5	590	1000		0	
Hg (dissolved)	2	0.01	0.01	0.022	0.0214	0.022	0.05		0	
Ni (dissolved)	2	1	1	1	1	1	20		0	
Pb (dissolved)	2	1	1	2	1.95	2	7.2		0	
Sn (dissolved)	2	1	1	2.2	2.14	2.2	25		0	
V (dissolved)	2	1	1	2.6	2.52	2.6	20		0	
Zn (total)	2	1	3.5	3.8	3.785	3.8	50		0	
Cyanide (free)	2	5	5	5	5	5	1		2	EQS > LoD.
Chloride (Cl-)	2	1000	2500	2700	2690	2700	250000		0	
Fluoride (F-)	2	50	160	250	245.5	250	5000		0	
Sulfate (SO42-)	2	1000	3600	6800	6640	6800	400000		0	
pH (min.) (su)	2	0.1	8.2	8	8.19	8	6		0	Max & Min interchanged to compare min. value.
pH (max.) (su)	2	0.1	8	8.2	8.19	8.2	8.5		0	
Anthracene	2	0.01	0.01	0.01	0.01	0.01	0.1		0	
Benzo(a)pyrene	2	0.01	6.7	8.9	8.79	8.9	0.05		2	
PAH sum of										
benzo(b)fluoranthene	-	0.00	0.00	4.54	4 4	4.54	0.00			
benzo(k)fluoranthene	2	0.02	0.02	1.21	1.1505	1.21	0.03		1	EQS > LoD.
PAH sum of										
benzo(ghi)perylene										
indeno(1,2,3-cd)pyrene	2	0.02	0.02	0.02	0.02	0.02	0.002		2	
Fluoranthene	2	0.01	0.01	0.6	0.5705	0.6	0.1		1	
Naphthalene	2	0.01	2.6	3.3	3.265	3.3	2.4		2	
Phenol	2	0.2	0.2	0.2	0.2	0.2	7.7		0	
2,4,6-Trichlorophenol	2	0.2	0.2	0.2	0.2	0.2			0	
2-Chlorophenol	2	0.2	0.2	0.2	0.2	0.2	50		0	
2,4-Dichlorophenol	2	0.2	0.2	0.2	0.2	0.2	20		0	
4-Chloro, 3-methylphenol	2	0.2	0.2	0.2	0.2	0.2	40		0	

# Summary of Remedial Targets Methodology

RTM Level 1 - Soil Zone A	ssessment	- leachate	samples								
Water body receptor(s)	: Surface wat	er									
Secondary receptor(s)	: Aquatic eco	system									
Data set: Tip Area											
Client: Bovis, Barratt and Taylor Wimpey Homes											
Site	Site: Land at Bankside, Banbury										
Job no	: C12702		-								
Chemicals of Potential Concern		Summa	ary of Sample	Data		Value Being Compared to Target =	(Exceed	llity Target ed if Red ext)	Exceedi	amples ng Water 7 Target	Notes
(concentrations in µg/l)	No. of Samples	Limit of Detection	Minimum Value	Maximum Value	95-%ile Value	Maximum Value	Inland Waters EQS		Inland Waters EQS		
Pentachlorophenol	2	0.2	0.2	0.2	0.2	0.2	0.4		0		

RTM Level 1 - Soil Zone	Assessmer	nt - leachat									
Water body receptor(s)											
Secondary receptor(s)			e waler								
		area (combine	(h)								
		att and Taylo		mor							
		nkside, Banbu		lies							
	: C12702	ikside, Ballot	y								
000 110	. 012/02					1					
		<b>C</b>	ary of Sample	Dete		Value Being	Water Qua				Notes
Chemicals of Potential		Summa	ary of Sample	Compared to	(Exceeded if Red Text)		Exceeding Water		General comment: where more than one		
Concern			Target =	i ext)		Quality Target		LoD applies because several labs are			
(concentrations in µg/l)	No. of	Limit of	Minimum	Maximum	Inland		Inland		involved, the highest is guoted.		
	Samples	Detection	Value	Maximum Value	95-%ile Value	Value	Waters		Waters		involved, the highest is quoted.
							EQS		EQS		
Ag (dissolved)	2	0.5	0.5	0.5	0.5	0.5	0.05		2		EQS < LoD.
As (dissolved)	21	1	1	4	3	4	50		0		
B (dissolved)	21	50	10	66	65	66	2000		0		
Cd (dissolved)	21	0.5	0.08	0.5	0.5	0.5	0.08		19		EQS < higher LoD.
Co (dissolved)	2	1	1	1	1	1	3		0		
Cr (VI) (dissolved)	12	2	1	2	2	2	3.4		0		
Cr (III) (dissolved)	15	2	1	8	5.9	8	4.7		2		
Cu (dissolved)	21	5	1	100	28	100	1		20		EQS < higher LoD, but some values are > EQS.
Fe (dissolved)	21	20	1	1200	220	1200	1000		1		
Hg (dissolved)	20	0.2	0.01	0.2	0.2	0.2	0.05		16		EQS <higher lod.<="" td=""></higher>
Ni (dissolved)	21	2	1	20	3.2	20	20		0		
Pb (dissolved)	2	1	1	1	1	1	7.2		0		
Sn (dissolved)	2	1	1	1	1	1	25		0		
V (dissolved)	2	1	1	1.8	1.76	1.8	20		0		
Zn (total)	21	5	1	27	10	27	8		4		EQS < LoD.
Cyanide (free)	12	5	5	5	5	5	1		12		EQS < LOD.
Chloride (Cl-)	2	1000	1400	2900	2825	2900	250000		0		
Fluoride (F-)	2	50	340	620	606	620	1000		0		
Sulfate (SO42-)	2	1000	1600	6700	6445	6700	400000		0		Max & Min interchanged to compare min. value.
pH (min.) (su)	12 12	0.1	8.5 7.1	7.1 8.5	8.28 8.28	7.1 8.5	6 8.5		0		iviax & with interchanged to compare min. Value.
pH (max.) (su)	12	0.1	0.01	8.5 0.036	8.28 0.02995	0.036	8.5 0.1		0		
Anthracene	12	0.01	0.01	7.1	6.055	7.1	0.1		2		
Benzo(a)pyrene PAH sum of	12	0.01	0.02	7.1	6.000	1.1	0.00		2		EQS < higher LoD.
benzo(b)fluoranthene											Lao Shighor LOD.
benzo(k)fluoranthene	12	0.04	0.02	0.04	0.04	0.04	0.03		10		
PAH sum of	12	0.04	0.02	0.04	0.04	0.04	0.03		10		EQS < LoD.
benzo(ghi)perylene											
indeno(1,2,3-cd)pyrene	10	0.04	0.00	0.04	0.04	0.04	0.000		40		
	12	0.04	0.02	0.04	0.04	0.04	0.002		12		
Fluoranthene	12	0.02	0.01	1	0.4951	1	0.1		1		
Naphthalene	12	0.02	0.01	2.2	1.0835	2.2	2.4		0		
Phenol	12	0.5	0.2	0.5	0.5	0.5	7.7		0		

# **Summary of Remedial Targets Methodology**

# Notes to Remedial Targets Methodology Table(s)

- 1 Data from the Environment Agency Chemical Standards web site at http://87.84.223.229/ChemicalStandards/Home.aspx
- 2 Substances in **bold** are defined in 2008/105/EC Annex II as priority substances in the field of water policy and those in **bold italic** as priority hazardous substances.
- 3 EQS for inland waters applies to freshwater rivers, lakes etc. EQS for other waters refers to marine and transitional (eg estuarine) waters.
- 4 Inland waters EQS for Cd, Cu and Zn depend on water hardness (mg/l as CaCO₃). Where applicable, water hardness is measured, otherwise it has been estimated by reference to the map at http://dwi.defra.gov.uk/consumers/advice-leaflets/hardness_map.pdf. If hardness cannot be determined, a worst case is assumed by setting it to 10mg/l.

# Appendix F

WASTE MANAGEMENT

# WASTE MANAGEMENT

## Establishing if Substances are Wastes

Any material excavated on site may be classified as waste and it is the responsibility of the holder of a material to form their own view on whether or not it is waste. This includes determining when waste that has been treated in some way can cease to be classed as waste for a particular purpose.

One of the ways this can be achieved is set out in the Development Industry Code of Practice (CoP) (CL:AIRE, March 2011). This builds on the Environment Agency guidance document *Definition of waste: developing greenfield and brownfield sites* (2006).

The handling, re-use or disposal of waste is regulated by the Agency. The Agency will take into account the use of the CoP in deciding whether to regulate materials as waste. If materials are dealt with in accordance with the CoP, the Agency considers that those materials are unlikely to be waste at the point when they are to be used for the purpose of land development. This may be because the materials were never discarded in the first place, or because they have been submitted to a recovery operation and have been completely recovered so that they have ceased to be waste.

Good practice has three basic elements:

- 1. Ensuring that an adequate Materials Management Plan (MMP) is in place, covering the use of materials on a specific site.
- 2. Ensuring that the MMP is based on an appropriate risk assessment, that underpins the Remediation Strategy or Design Statement, concluding that the objectives of preventing harm to human health and pollution of the environment will be met if materials are used in the proposed manner.
- 3. Ensuring that materials are actually treated and used as set out in the MMP and that this is subsequently demonstrated in a Verification Report.

To confirm that Steps 1 and 2 have been taken, a Qualified Person must review the relevant documents and provide a Declaration before excavation work commences on a particular site. When the Declaration is provided to the Agency, demonstrating that the materials are to be dealt within accordance with the MMP and risk assessment, the Agency will take the view that the materials on the site where they are to be used will not be waste.

However, if it turns out that materials were not used in accordance with the MMP and risk assessment, or if it is discovered that materials are not suitable for use, are used or planned to be used in excessive quantities, or are likely to cause harm to human health or pollution of the environment, the Agency may conclude that those materials have been discarded and are waste. In order to show how materials have been treated and used, a Verification Report must be prepared at the conclusion of works and, if requested, provided to the Agency. Completion of a Verification Report will not prevent consideration of the above matters by the Agency.

The CoP applies to both uncontaminated and contaminated material from anthropogenic and natural sources excavated:

- for use on the site from which it has been excavated, either without treatment or after onsite treatment, as part of the development of that land (Site of Origin scenario);
- for use directly without treatment at another development site subject to the material meeting certain requirements (Direct Transfer scenario);
- for use in the development of land other than the site from which the material has been excavated, following treatment at an authorised Hub site including a fixed Soil Treatment Facility (Cluster Project scenario); or
- a combination of the above.

The need to distinguish between "contaminated" and "uncontaminated" soils is no longer considered necessary. The Agency accepts that these are self-defining terms on a site specific basis having regard to the risk assessment (e.g. some soil may not be considered contaminated for a given land use, but would be for a more sensitive land use, on the same site).

The fact that the material has to be treated indicates that it is a waste, i.e. it is not suitable for use until it is treated.

"Development" also includes redevelopment, remediation and regrading of a site. The CoP therefore applies not only to development carried out under the development control regime, but also to remediation activities which may occur outside of that regime, e.g. remediation as a direct result of a spillage or leak on an industrial site or at the surrender stage of a permit. Land development or remediation does not include landfilling of waste or other waste disposal operations.

There is no single factor that can be used to determine if something is a waste or when it ceases to be waste. However, in the context of excavated materials used on sites undergoing development, the following factors are considered to be of particular relevance.

- Factor 1: protection of human health and the environment Compliance with the Water Framework Directive. If the use of the material will create unacceptable risk, it is likely to be waste.
- Factor 2: suitability for use, without any further treatment Suitability for use means that a material must be suitable for its intended purpose in all respects. In particular, both its chemical and geotechnical properties have to be demonstrated to be suitable, and the relevant specification for its use must be met. Suitability of use also includes consideration of the effect that the material may have on the environment.

Certain excavated materials may be suitable for their intended use in the proposed development without any treatment at all. If they are used in that way those materials are unlikely to be waste. For example some materials may be assessed as being suitable for direct use, e.g. engineered backfill beneath cover layers, capping layers, buildings and hard standing or for site regrading. Use for the purposes of reclamation, restoration, landscaping or improvement of land may fall within this category. Landfilling or disposal

#### does not.

Other materials may not have the required characteristics for use without first being treated. If treatment is needed in order to make the material ready for use the materials will be waste, but may cease to be waste once treated so as to be suitable for use. This treatment may be biological, chemical, physical or any combination of these and will need to be carried out under an appropriate authorisation.

Some materials, although they do not require treatment to make them suitable for use, may nonetheless be regraded or compacted before or during their use as part of the development of a site. This regrading or compacting does not prevent the material being regarded as a non-waste.

 Factor 3: certainty of use - The holder of the material must be able to demonstrate that the material will actually be used and that the use is not just a probability, but a certainty. For example, if materials are stockpiled with no pre-defined destination and use, they will be waste.

In the process of site development surplus material may be generated that cannot be used either directly or after treatment. For example, the material may not conform to the required specification following treatment and in such a case the material would remain a waste.

There may be unexpected arisings on a development site that were not picked up within the site investigation works. Any out of specification materials will be waste and will need to be disposed of or recovered in the proper manner and in accordance with waste legislation.

• Factor 4: quantity of material - Materials should be used in the quantities necessary for that use, and no more. The use of an excessive amount of material will indicate that it is being disposed of and is waste.

The production of a MMP will help to ensure that the above matters are considered and a correct determination is made in relation to the nature of the materials.

In order to demonstrate that the factors described above have been satisfied, an MMP has to be produced relating to the use of the materials, which includes a tracking system and contingency arrangements. The MMP accompanies the Remediation Strategy or Design Statement, which has been derived using risk assessment. A Verification Plan also needs to form a part of the MMP.

Upon completion of these documents, a Qualified Person needs to sign a Declaration. Once the development has been completed in accordance with the MMP, a Verification Report must be completed that demonstrates that the materials have been located in the correct place within the development or dealt with appropriately.

Further details can be found in the CoP.

# Key Legal Requirements

If the material is considered to be waste then the legislation will apply up to the point that it ceases to be waste.

**Duty of Care:** It is necessary to ensure all waste is handled, recovered and disposed of responsibly, and that the waste is only handled by individuals, companies or groups that are authorised to deal with it. For example, waste can only be collected by registered carriers or transporters. Regular checks must be made on the destination of all wastes leaving site to ensure they are only being taken to an appropriately authorised waste management facility. Records (Waste Transfer Notes) must be kept of all wastes received or transferred.

Characteristics of waste received from a third party must be checked its to ensure that companies are licensed or have an exemption under which they can receive it and that it complies with the classification set out in the Waste Transfer Notes.

Waste Carrier or Transporter: Registration is require to transport waste.

**Environmental Permits:** It is normally an offence to undertake waste disposal or recovery operations, which are regulated by the Agency, without being in possession of an appropriate Environmental Permit, unless it is material that is "uncontaminated soil and other naturally occurring material excavated in the course of construction activities where it is certain that the material will be used for the purposes of construction in its natural state on the site from which it was excavated", which is excluded from waste regulation by the Waste Framework Directive.

As an alternative to using the Industry CoP in excavating and re-using materials there are a number of options:

- Waste Exemption mainly for small volumes of non-hazardous waste, recovery only;
- Standard Rules Environmental Permit which replaces the traditionally used Waste Exemptions Paragraphs 9 and 19, but can take several months to obtain;
- Bespoke Environmental Permit applicable to greater volumes and more waste streams than Standard Rules, but can take several months to obtain; or
- WRAP Aggregates Quality Protocol allows for inert aggregate waste to be recovered and used at any site subject to meeting set standards.

# Waste Classification

With respect to the possible waste streams from a site, it is recommended that a phased approach is implemented. In the first instance, the groundwork's contractor or specialist remediation contractor appointed by the developer should approach the landfill site with the available chemical data and seek a waste characterisation. Should the waste be classified as hazardous, it would be necessary to undertake the Waste Acceptance Criteria (WAC) testing to determine whether the receiving landfill could accept the hazardous waste. This would require additional soil sampling and chemical testing.

The two stages are explained below.

# Waste Characterisation

All wastes going to landfill must be classified as 'inert', 'non-hazardous' or 'hazardous'. There is a sub-category of hazardous waste known as 'stable non-reactive hazardous waste'. Individual landfill sites have permits to take these classes of waste. Hazardous and non-hazardous wastes cannot be disposed of at the same site, apart from stable non-reactive hazardous waste which can go to specially constructed cells in certain non-hazardous landfill sites.

Contaminated soil is a 'mirror entry' in the Consolidated European Waste Catalogue, and is not necessarily a hazardous waste. It is only classified as hazardous if it contains dangerous substances above certain threshold concentrations. The Environment Agency Briefing Note on Hazardous Waste and Contaminated Soil (V.1 July 2004) suggests that waste holders should use the information collected as part of the contaminated land risk assessment to inform decisions as to the concentrations that might reasonably be expected to be present in the contaminated soil, given the past and current uses of the site.

The waste must be assessed against all the appropriate hazards in accordance with the Environment Agency Technical Guidance WM2. This makes certain worst case assumptions about the chemical composition if specific compounds are not analysed for.

# Waste Acceptance Criteria

Waste classified as hazardous must be subject to WAC testing to determine if it can go to a hazardous landfill site. The WAC are a list of limit values for certain parameters obtained from standard leaching tests and total content tests. If the limit values are exceeded, the waste is not suitable for disposal at that class of landfill site and alternative disposal methods have to be found. Maximum permissible limit values are determined by the EU (part of what is known as 'full waste acceptance criteria') but individual landfills may have more stringent values to take into account the environmental setting, liner system or additional nature of specific waste streams.

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