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Newport City Council
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Newport
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Attention: Robert Hester

CERTIFICATE OF ANALYSIS

Date of report Generation: 13 February 2020
Customer: Newport City Council
Sample Delivery Group (SDG): 200205-66
Your Reference: Jan Leachate
Location: Docks Way
Report No: 541170

We received 2 samples on Wednesday February 05, 2020 and 2 of these samples were scheduled for analysis which was completed on Thursday February 13, 2020. Accredited laboratory tests are defined within the report, but opinions, interpretations and on-site data expressed herein are outside the scope of ISO 17025 accreditation.

Should this report require incorporation into client reports, it must be used in its entirety and not simply with the data sections alone.

Chemical testing (unless subcontracted) performed at ALS Environmental Hawarden (Method codes TM) or ALS Environmental Aberdeen (Method codes S).

All sample data is provided by the customer. The reported results relate to the sample supplied, and on the basis that this data is correct.

Incorrect sampling dates and/or sample information will affect the validity of results.

The customer is not permitted to reproduce this report except in full without the approval of the laboratory.

Approved By:

Sonia McWhan

Operations Manager





CERTIFICATE OF ANALYSIS

Validated

SDG:	200205-66	Client Reference:	Jan Leachate	Report Number:	541170
Location:	Docks Way	Order Number:	700145760	Superseded Report:	

Received Sample Overview

Lab Sample No(s)	Customer Sample Ref.	AGS Ref.	Depth (m)	Sampled Date
21624238	C3		0.00 - 0.00	05/02/2020
21624251	C4		0.00 - 0.00	05/02/2020

Maximum Sample/Coolbox Temperature (°C) : 9.8

ISO5667-3 Water quality - Sampling - Part3 -

During Transportation samples shall be stored in a cooling device capable of maintaining a temperature of (5±3)°C.

ALS have data which show that a cool box with 4 frozen icepacks is capable of maintaining pre-chilled samples at a temperature of (5±3)°C for a period of up to 24hrs.

Only received samples which have had analysis scheduled will be shown on the following pages.



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Results Legend		Customer Sample Ref.	C3	C4			
# ISO17025 accredited. M mCERTS accredited. aq Aqueous / settled sample. diss.filt Dissolved / filtered sample. tot.unfilt Total / unfiltered sample. * Subcontracted - refer to subcontractor report for accreditation status. ** % recovery of the surrogate standard to check the efficiency of the method. The results of individual compounds within samples aren't corrected for the recovery. (F) Trigger breach confirmed 1.3.6@ Sample deviation (see appendix)		Depth (m) Sample Type Date Sampled Sampled Time Date Received SDG Ref Lab Sample No.(s) AGS Reference	0.00 - 0.00 Land Leachate (LE) 05/02/2020 . 05/02/2020 200205-66 21624238	0.00 - 0.00 Land Leachate (LE) 05/02/2020 . 05/02/2020 200205-66 21624251			
Component	LOD/Units	Method					
Ionic balance		Calulation	2.49	4.72			
	% Diff						
Alkalinity, Total as CaCO3 (diss.filt)	<2 mg/l	TM043	8740	2600			
Alkalinity, Bicarbonate as CaCO3 (diss.filt)	<2 mg/l	TM043	8740	2600			
BOD, unfiltered	<1 mg/l	TM045	406	196	#	#	
Organic Carbon, Total	<3 mg/l	TM090	2090	446			
Ammoniacal Nitrogen as N	<0.2 mg/l	TM099	1920	312			
Sulphide	<0.01 mg/l	TM101	0.506	0.177		2	
COD, unfiltered	<7 mg/l	TM107	7980	1420	#	#	
Conductivity @ 20 deg.C	<0.005 mS/cm	TM120	20.7	10.9	#	#	
Arsenic (diss.filt)	<0.5 µg/l	TM152	80.4	164	2 #	#	
Boron (diss.filt)	<10 µg/l	TM152	27300	8390	2 #	#	
Cadmium (diss.filt)	<0.08 µg/l	TM152	<0.8	<0.08	2 #	#	
Chromium (diss.filt)	<1 µg/l	TM152	904	79.8	2 #	#	
Copper (diss.filt)	<0.3 µg/l	TM152	<3	2.92	2 #	#	
Lead (diss.filt)	<0.2 µg/l	TM152	6.69	0.894	2 #	#	
Manganese (diss.filt)	<3 µg/l	TM152	354	1780	2 #	#	
Nickel (diss.filt)	<0.4 µg/l	TM152	247	95.5	2 #	#	
Selenium (diss.filt)	<1 µg/l	TM152	<10	3.02	2 #	#	
Zinc (diss.filt)	<1 µg/l	TM152	78.7	18.7	2 #	#	
Sodium (Dis.Filt)	<0.076 mg/l	TM152	2360	1370	2 #	#	
Magnesium (Dis.Filt)	<0.036 mg/l	TM152	214	252	2 #	#	
Potassium (Dis.Filt)	<0.2 mg/l	TM152	632	296	2 #	#	
Calcium (Dis.Filt)	<0.2 mg/l	TM152	71	697	2 #	#	
Iron (Dis.Filt)	<0.019 mg/l	TM152	2.46	0.167	2 #	#	
Hardness, Total as CaCO3	<0.65 mg/l	TM152	1060	2780	2		
EPH Range >C10 - C40 (aq)	<100 µg/l	TM172	3050	1500			
Nitrite as NO2	<0.05 mg/l	TM184	<0.05	<0.05			
Phosphate (Ortho as PO4)	<0.05 mg/l	TM184	41.2	1.94			
Sulphate	<2 mg/l	TM184		111			
Chloride	<2 mg/l	TM184		2760			
Nitrate as NO3	<0.3 mg/l	TM184		<0.3			
Total Oxidised Nitrogen as N	<0.1 mg/l	TM184		<0.1			



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Table of Results - Appendix

Method No	Reference	Description
Calculation		
TM043	Method 2320B, AWWA/APHA, 20th Ed., 1999 / BS 2690: Part109 1984	Determination of alkalinity in aqueous samples
TM045	MEWAM BOD5 2nd Ed.HMSO 1988 / Method 5210B, AWWA/APHA, 20th Ed., 1999; SCA Blue Book 130	Determination of BOD5 (ATU) Filtered by Oxygen Meter on liquids
TM090	Method 5310, AWWA/APHA, 20th Ed., 1999 / Modified: US EPA Method 415.1 & 9060	Determination of Total Organic Carbon/Total Inorganic Carbon in Water and Waste Water
TM099	BS 2690: Part 7:1968 / BS 6068: Part2.11:1984	Determination of Ammonium in Water Samples using the Kone Analyser
TM101	Method 4500B & C, AWWA/APHA, 20th Ed., 1999	Determination of Sulphide in soil and water samples using the Kone Analyser
TM107	ISO 6060-1989	Determination of Chemical Oxygen Demand using COD Dr Lange Kit
TM120	Method 2510B, AWWA/APHA, 20th Ed., 1999 / BS 2690: Part 9:1970	Determination of Electrical Conductivity using a Conductivity Meter
TM152	Method 3125B, AWWA/APHA, 20th Ed., 1999	Analysis of Aqueous Samples by ICP-MS
TM172	Analysis of Petroleum Hydrocarbons in Environmental Media – Total Petroleum Hydrocarbon Criteria	EPH in Waters
TM184	EPA Methods 325.1 & 325.2,	The Determination of Anions in Aqueous Matrices using the Kone Spectrophotometric Analysers
TM208	Modified: US EPA Method 8260b & 624	Determination of Volatile Organic Compounds by Headspace / GC-MS in Waters
TM226	In-House Method	Determination of Anions in Waters using Ion Chromatography
TM227	Standard methods for the examination of waters and wastewaters 20th Edition, AWWA/APHA Method 4500.	Determination of Total Cyanide, Free (Easily Liberatable) Cyanide and Thiocyanate
TM256	The measurement of Electrical Conductivity and the Laboratory determination of pH Value of Natural, Treated and Wastewaters. HMSO, 1978. ISBN 011 751428 4.	Determination of pH in Water and Leachate using the GLpH pH Meter

NA = not applicable.

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Test Completion Dates

Table with 3 columns: Lab Sample No(s), Customer Sample Ref., AGS Ref., Depth, Type. Rows include sample numbers 21624238 and 21624251, customer refs C3 and C4, AGS Ref., Depth 0.00 - 0.00, and Type Land Leachate.

Main test results table with 3 columns: Test Name, 21624238, 21624251. Rows include Alkalinity Filtered as CaCO3, Ammoniacal Nitrogen, Anions by ion Chromatography, Anions by Kone (w), BOD True Total, COD Unfiltered, Conductivity (at 20 deg.C), Cyanide Comp/Free/Total/Thiocyanate, Dissolved Metals by ICP-MS, EPH (DRO) (C10-C40) Aqueous (W), Ionic Balance, Nitrite by Kone (w), pH Value, Phosphate by Kone (w), Sulphide, Total Organic and Inorganic Carbon, and VOC MS (W).



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Chromatogram

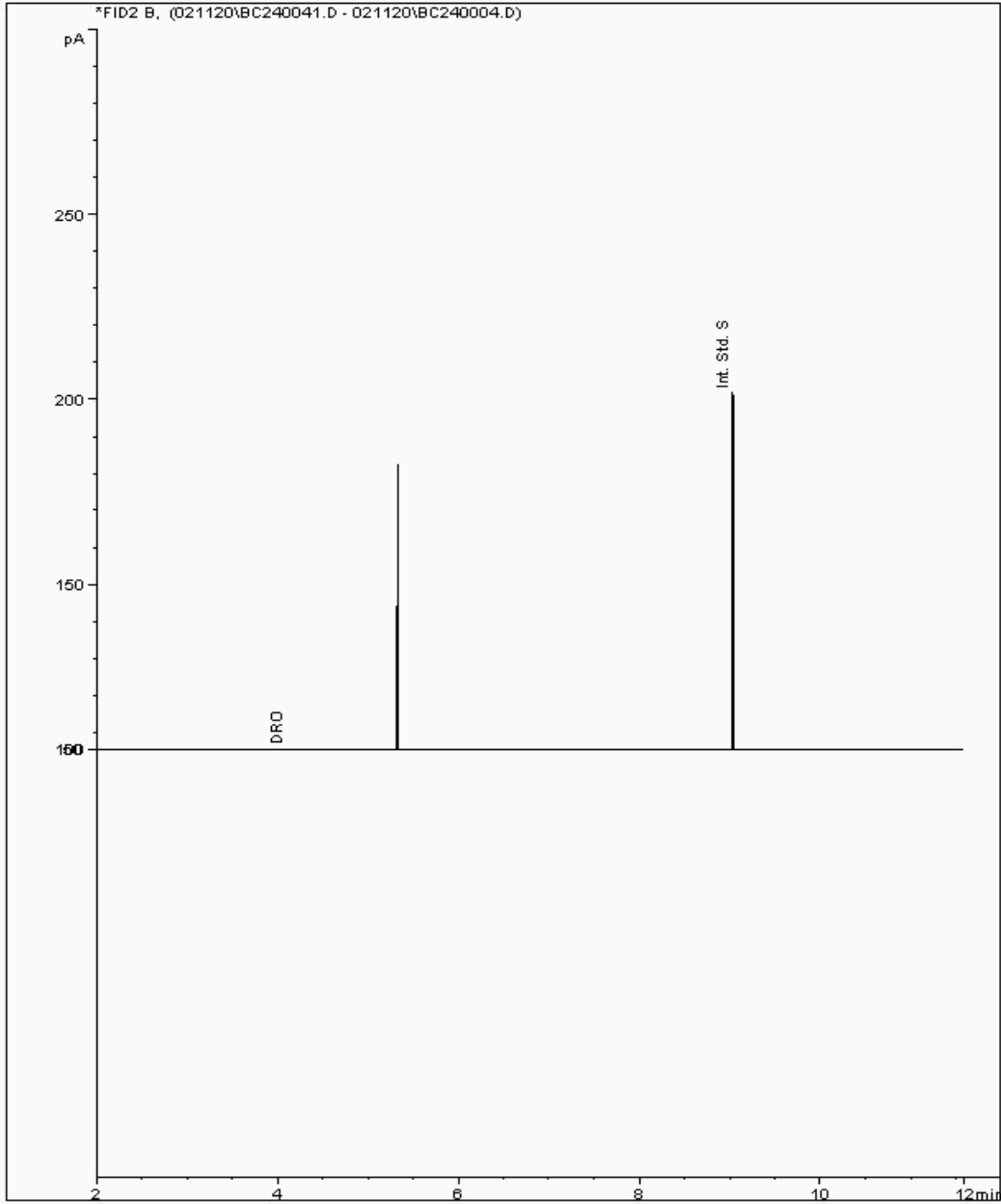
Analysis: EPH (DRO) (C10-C40) Aqueous (W)

Sample No : 21624719
Sample ID : C3

Depth : 0.00 - 0.00

EPH Range Organics (C10 - C40)

Sample Identity: 20309769-
Date Acquired : 11/02/2020 23:29:44 PM
Units : mg/l





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Chromatogram

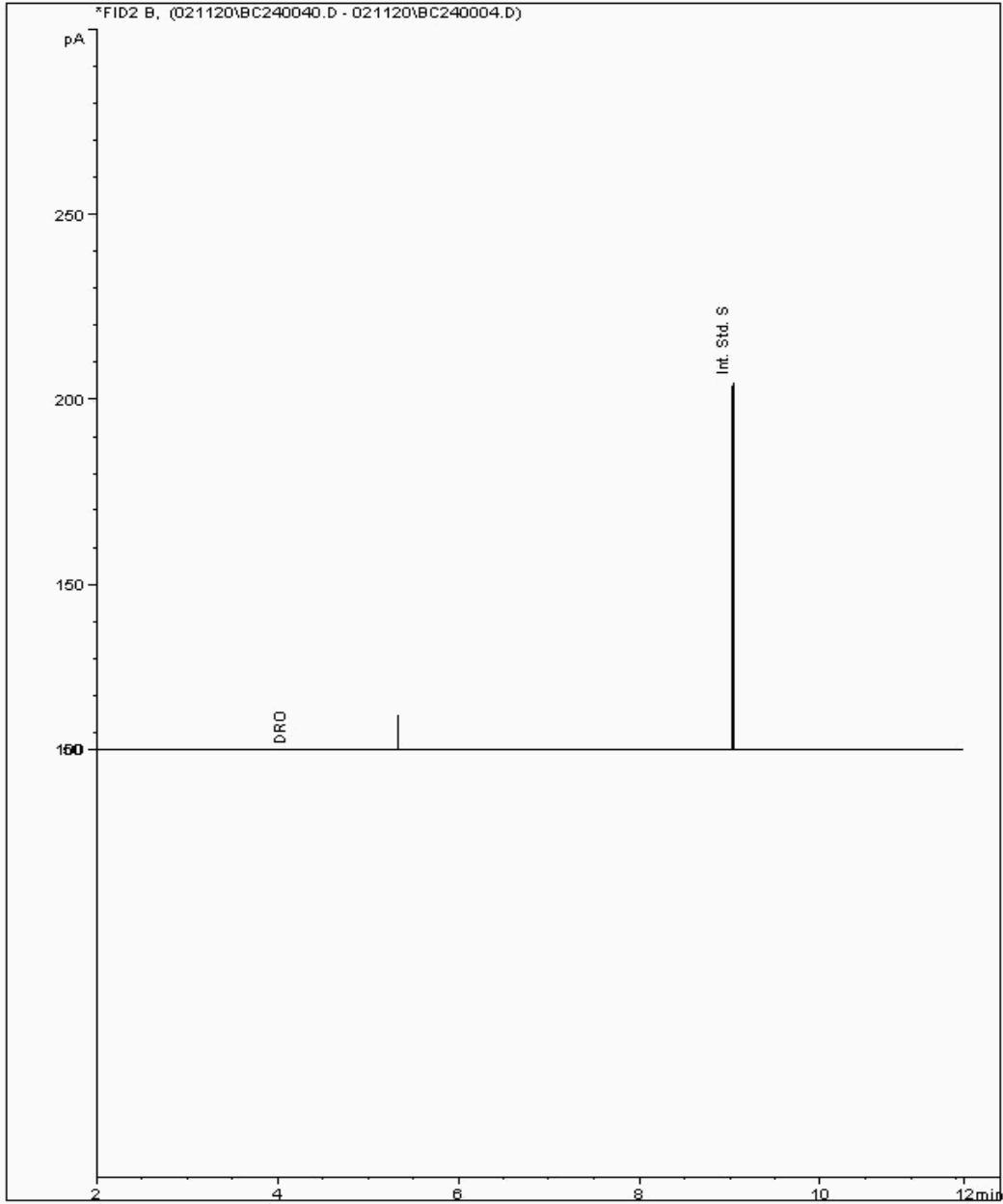
Analysis: EPH (DRO) (C10-C40) Aqueous (W)

Sample No : 21624723
Sample ID : C4

Depth : 0.00 - 0.00

EPH Range Organics (C10 - C40)

Sample Identity: 20309791-
Date Acquired : 11/02/2020 23:06:08 PM
Units : mg/l





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Appendix

General

1. Results are expressed on a dry weight basis (dried at 35°C) for all soil analyses except for the following: NRA and CEN Leach tests, flash point LOI, pH, ammonium as NH4 by the BRE method, VOC TICs and SVOC TICs.

2. If sufficient sample is received a sub sample will be retained free of charge for 30 days after analysis is completed (e-mailed) for all sample types unless the sample is destroyed on testing. The prepared soil sub sample that is analysed for asbestos will be retained for a period of 6 months after the analysis date. All bulk samples will be retained for a period of 6 months after the analysis date. All samples received and not scheduled will be disposed of one month after the date of receipt unless we are instructed to the contrary. Once the initial period has expired, a storage charge will be applied for each month or part thereof until the client cancels the request for sample storage. ALS reserve the right to charge for samples received and stored but not analysed.

3. With respect to turnaround, we will always endeavour to meet client requirements wherever possible, but turnaround times cannot be absolutely guaranteed due to so many variables beyond our control.

4. We take responsibility for any test performed by sub-contractors (marked with an asterisk). We endeavour to use UKAS/MCERTS Accredited Laboratories, who either complete a quality questionnaire or are audited by ourselves. For some determinands there are no UKAS/MCERTS Accredited Laboratories, in this instance a laboratory with a known track record will be utilised.

5. If no separate volatile sample is supplied by the client, or if a headspace or sediment is present in the volatile sample, the integrity of the data may be compromised. This will be flagged up as an invalid VOC on the test schedule and the result marked as deviating on the test certificate.

6. NDP - No determination possible due to insufficient/unsuitable sample.

7. Results relate only to the items tested.

8. LoDs (Limit of Detection) for wet tests reported on a dry weight basis are not corrected for moisture content.

9. **Surrogate recoveries** - Surrogates are added to your sample to monitor recovery of the test requested. A % recovery is reported, results are not corrected for the recovery measured. Typical recoveries for organics tests are 70-130%. Recoveries in soils are affected by organic rich or clay rich matrices. Waters can be affected by remediation fluids or high amounts of sediment. Test results are only ever reported if all of the associated quality checks pass; it is assumed that all recoveries outside of the values above are due to matrix affect.

10. Stones/debris are not routinely removed. We always endeavour to take a representative sub sample from the received sample.

11. In certain circumstances the method detection limit may be elevated due to the sample being outside the calibration range. Other factors that may contribute to this include possible interferences. In both cases the sample would be diluted which would cause the method detection limit to be raised.

12. Mercury results quoted on soils will not include volatile mercury as the analysis is performed on a dried and crushed sample.

13. For leachate preparations other than Zero Headspace Extraction (ZHE) volatile loss may occur.

14. For the BSEN 12457-3 two batch process to allow the cumulative release to be calculated, the volume of the leachate produced is measured and filtered for all tests. We therefore cannot carry out any unfiltered analysis. The tests affected include volatiles GCFID/GCMS and all subcontracted analysis.

15. Analysis and identification of specific compounds using GCFID is by retention time only, and we routinely calibrate and quantify for benzene, toluene, ethylbenzenes and xylenes (BTEX). For total volatiles in the C5-C12 range, the total area of the chromatogram is integrated and expressed as ug/kg or ug/l. Although this analysis is commonly used for the quantification of gasoline range organics (GRO), the system will also detect other compounds such as chlorinated solvents, and this may lead to a falsely high result with respect to hydrocarbons only. It is not possible to specifically identify these non-hydrocarbons, as standards are not routinely run for any other compounds, and for more definitive identification, volatiles by GCMS should be utilised.

16. We are accredited to MCERTS for sand, clay and loam/topsoil, or any of these materials - whether these are derived from naturally occurring soil profiles, or from fill/made ground, as long as these materials constitute the major part of the sample. Other coarse granular material such as concrete, gravel and brick are not accredited if they comprise the major part of the sample.

17. **Tentatively Identified Compounds (TICs)** are non-target peaks in VOC and SVOC analysis. All non-target peaks detected with a concentration above the LoD are subjected to a mass spectral library search. Non-target peaks with a library search confidence of >75% are reported based on the best mass spectral library match. When a non-target peak with a library search confidence of <75% is detected it is reported as "mixed hydrocarbons". Non-target compounds identified from the scan data are semi-quantified relative to one of the deuterated internal standards, under the same chromatographic conditions as the target compounds. This result is reported as a semi-quantitative value and reported as Tentatively Identified Compounds (TICs). TICs are outside the scope of UKAS accreditation and are not moisture corrected.

18. Sample Deviations

If a sample is classed as deviated then the associated results may be compromised.

1	Container with Headspace provided for volatiles analysis
2	Incorrect container received
3	Deviation from method
§	Sampled on date not provided
◆	Sample holding time exceeded in laboratory
@	Sample holding time exceeded due to late arrival of instructions or samples

19. Asbestos

When requested, the individual sub sample scheduled will be analysed in house for the presence of asbestos fibres and asbestos containing material by our documented in house method TM048 based on HSG 248 (2005), which is accredited to ISO17025. If a specific asbestos fibre type is not found this will be reported as "Not detected". If no asbestos fibre types are found all will be reported as "Not detected" and the sub sample analysed deemed to be clear of asbestos. If an asbestos fibre type is found it will be reported as detected (for each fibre type found). Testing can be carried out on asbestos positive samples, but, due to Health and Safety considerations, may be replaced by alternative tests or reported as No Determination Possible (NDP). The quantity of

Identification of Asbestos in Bulk Materials & Soils

The results for identification of asbestos in bulk materials are obtained from supplied bulk materials which have been examined to determine the presence of asbestos fibres using ALS (Hawarden) in-house method of transmitted/polarised light microscopy and central stop dispersion staining, based on HSG 248 (2005).

The results for identification of asbestos in soils are obtained from a homogenised sub sample which has been examined to determine the presence of asbestos fibres using ALS (Hawarden) in-house method of transmitted/polarised light microscopy and central stop dispersion staining, based on HSG 248 (2005).

Asbestos Type	Common Name
Chrysotile	White Asbestos
Amosite	Brown Asbestos
Crocidolite	Blue Asbestos
Fibrous Actinolite	-
Fibrous Anthophyllite	-
Fibrous Tremolite	-

Visual Estimation Of Fibre Content

Estimation of fibre content is not permitted as part of our UKAS accredited test other than: - Trace - Where only one or two asbestos fibres were identified.

Respirable Fibres

Respirable fibres are defined as fibres of <3 µm diameter, longer than 5 µm and with aspect ratios of at least 3:1 that can be inhaled into the lower regions of the lung and are generally acknowledged to be most important predictor of hazard and risk for cancers of the lung.

Standing Committee of Analysts, *The Quantification of Asbestos in Soil (2107)*.

Further guidance on typical asbestos fibre content of manufactured products can be found in HSG 264.

The identification of asbestos containing materials and soils falls within our schedule of tests for which we hold UKAS accreditation, however opinions, interpretations and all other information contained in the report are outside the scope of UKAS accreditation.