GROUNDWATER ANALYTICAL

September 22, 2005

Groundwater Analytical, Inc. P.O. Box 1200 228 Main Street Buzzards Bay, MA 02532

Telephone (508) 759-4441 FAX (508) 759-4475 www.groundwateranalytical.com

Mr. Kevin Trainer Geolnsight, Inc. 5 Lan Drive Second Floor Westford. MA 01886

LABORATORY REPORT

Project:

Buzzards Bay/3871-002

Lab ID:

87247

Received:

08-30-05

Dear Kevin:

Enclosed are the analytical results for the above referenced project. The project was processed for Standard turnaround.

This letter authorizes the release of the analytical results, and should be considered a part of this report. This report contains a sample receipt report detailing the samples received, a project narrative indicating project changes and non-conformances, a quality control report, and a statement of our state certifications.

The analytical results contained in this report meet all applicable NELAC standards, except as may be specifically noted, or described in the project narrative. This report may only be used or reproduced in its entirety.

I attest under the pains and penalties of perjury that, based upon my inquiry of those individuals immediately responsible for obtaining the information, the material contained in this report is, to the best of my knowledge and belief, accurate and complete.

Should you have any questions concerning this report, please do not hesitate to contact me.

Sincerely,

Eric H. Jensey

Operations Manager

EHJ/kal

Enclosures



Sample Receipt Report

Project: Client:

Buzzards Bay/3871-002

Geolnsight, Inc.

Lab ID: 87247

Delivery: Hand

Airbill: n/a

Lab Receipt: 08-30-05

Temperature: 4.8'C

Chain of Custody: Present

Custody Seal(s): n/a

Lab ID	Field ID		Matrix	Sampled	Method	-1745.24.0 (F) (F) (F) -14.2 (Faul (MEDI) (F)			Notes
87247-1	W2A02-82905-01		Soil	8/29/05 11:30	MA DEP EPH	with PAHs by	8270C-Mod S	М	
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship		
C613445	120 mL Amber Glass	Proline	BX17622	None	n/a	n/a	n/a		

Lab ID	Field 1D		Matrix	Sampled	Method				Notes
87247-2	W2A02-82905-02		Soil	8/29/05 11:32	MA DEP EPH	with PAHs by	8270C-Mod S	IM	
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship		
C613447	120 mL Amber Glass	Proline	BX17622	None	n/a	n/a	n/a		



Data Certification

Project: Client: Buzzards Bay/3871-002

Geolnsight, Inc.

Lab ID:

87247

Received:

08-30-05 20:21

		ALCO 100 A CONTRACTOR AND ADDRESS OF THE ADDRESS OF		(1) (1) (1) (1) (1) (1) (1) (1) (1) (1)	***								
	And Andrews		MA DE	P Compen	dium o	f Analyti	cal Met	200 A CO. C.				Sa. 74	
Proj	ect Location:	n/a						MA	DEP RT	N:	r	n/a	
This	Form provides	certifications for the	e followir	g data set:									
MA	DEP EPH:	87247-01,-02											
Sam	ple Matrices:	Groundwater	() Soi	/Sediment	(X)	Drinking	2 Water	()	Oth	er	()		
	SW-846	8260B	()	8151A	()		8330	()	6010		()	7470A/1/	A (
Met	hods Used	8270C	()	8081A	()		VPH	()	60.		()	9012A	
55/2012/19/201	ecified in MA DEP pendium of Analytical	8082	()	8021B	()		EPH	(X)	7000	S ³	()	Othe	r (
Metho		List Release Tracking SW-846 Method 90		4	Samuel Contract Contract	EP Physiolo	eically Av	allable Cv	anide (PAC)	Methr	id .		
checl	k all that apply)	3. S - SW-846 Method						anabic Cy	amue (i /ic)	rectific			
	An affirmative re	esponse to question	ns A, B, C	and D is r	equire	d for "Pre	esumpti	ve Certa	inty" stat	us.	60 Aug 27 Thurston	- 17 (19 <u>19 19 19 19 19 19 19 19 19 19 19 19 19 1</u>	
Α.	Were all sa	mples received by t	the labora	atory in a c	onditio	on consis	tent wit	h					
		ed on the Chain-of									Υ	'es	
	14/- !! 0	1/06		41	· · · · ·		., .			-		**************************************	******
В.		VQC procedures re this report followe						(S)					
		narrative QC data		-									
		guidelines?		,	r - r						Υ	'es	
	D	- f. (* 1 - 1 - 1 - 1 - 1 - 1	3. 12. (1:							+			
2.		alytical data incluc ptive Certainty," as											
		CAM VII A, Quality						3					
.		isition and Report									Υ	es	
				(5)		,				-			
). D.		<u>'H methods only:</u> \nodifications, as sp				od run w	/ithout				V	es	
											1	<u>es</u>	
	A response to qu	estions E and F bel	low is rec	urred for "	Presum	nptive Ce	rtainty"	status.					
		performance stan	dards and	l recomme	ndatio	ns for the	9						
	specified m	ethods achieved?				***					Υ	es	
.	Were results	for all analyte-list	compour	ids/elemen	ts for t	he specif	ied			No.			
	method(s) re					,					Y	es	
	All No answers a	are addressed in th	e attache	d Project !	Varrati	ve.						*	
							at he-		n m:/ ==				
		attest under the ponsible for obta				-				rsor	idi		
-	-	to the best of my	-										
	-	_	1	4.0				-					
iana	iture:	601				Position	. ,	Inaratio	ine Mana	go-			
115110	ILLUIC.	97t1				OSILION	. (peralio	ns Mana	gei			
	ed Name:	Eric H. Jensen				Date:		9-22-05					



Massachusetts DEP EPH Method Extractable Petroleum Hydrocarbons by GC/FID

Field ID:

W2A02-82905-01

Project:

Buzzards Bay/3871-002

Client:

Geolnsight, Inc.

Laboratory ID: Sampled:

87247-01 08-29-05 11:30 08-30-05 20:21

Received: Extracted:

09-12-05 14:00 Analyzed (AL): 09-22-05 05:08 09-22-05 05:52

Analyzed (AR): Analyst:

MM

Matrix:

Soil

120 mL Amber Glass Container:

Preservation:

Cool

QC Batch ID:

EP-2142-M

Instrument ID:

GC-9 Agilent 6890

Sample Weight:

15 g

Final Volume: % Solids:

1 mL 88

Aliphatic Dilution Factor: 1 Aromatic Dilution Factor: 1

EPH Ranges Co	oncentration Not	tes Units	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	33
n-C19 to n-C36 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	33
n-C11 to n-C22 Aromatic Hydrocarbons ^{† o}	BRL	mg/Kg	33
<u>Unadjusted</u> n-C11 to n-C22 Aromatic Hydrocarbons [†]	BRL	mg/Kg	33

QC Surrogate C	ompound	Spiked	Measured	Recovery	QC Limits_
Fractionation:	2-Fluorobiphenyl	2.9	2.2	76 %	40 - 140 %
	2-Bromonaphthalene	2.9	2.3	79 %	40 - 140 %
Extraction:	Chloro-octadecane	2.9	2.1	72 %	40 - 140 %
	ortho -Terphenyl	2.9	2.2	74 %	40 - 140 %

	2-Bromonaphthalene	2.9	2.3	<i>7</i> 9 %	40 - 140 %
Extraction:	Chloro-octadecane	2.9	2.1	72 %	40 - 140 %
	ortho -Terphenyl	2.9	2.2	74 %	40 - 140 %
			ONIOC C-4	101 - 41	

QA/QC Certification

- 1. Were all QA/QC procedures required by the method followed?
- 2. Were all performance/acceptance standards for the required QA/QC procedures achieved?

Yes Yes

3. Were any significant modifications made to the method, as specified in Section 11.3.1.1?

No

Method non-conformances indicated above are detailed below on this data report, or in the accompanying project narrative and project quality control report. Release of this data is authorized by the accompanying signed project cover letter. The accompanying cover letter, project narrative and quality control report are considered part of this data report.

Method Reference:

Method for the Determination of Extractable Petroleum Hydrocarbons, MA DEP (Revision 1.1, 2004). Sample extraction performed by microwave accelerated solvent extraction technique. Results are reported on a dry weight basis.

Report Notations:

- Indicates concentration, if any, is below reporting limit for analyte. Reporting limit is the lowest concentration that can be reliably quantified under routine laboratory operating conditions. Reporting limits are adjusted for sample size and dilution.
- Hydrocarbon range data excludes concentrations of any surrogate(s) and/or internal standards eluting in that range.
- n-C11 to n-C22 Aromatic Hydrocarbons range data excludes the method target analyte concentrations.



EPA Method 8270C (Modified) MA DEP EPH Polynuclear Aromatic Hydrocarbons by GC/MS-SIM

Field ID:

W2A02-82905-01

Buzzards Bay/3871-002

Matrix: Container: Soil

Project:

Geolnsight, Inc.

Preservation:

120 mL Amber Glass Cool

.Client:

87247-01

QC Batch ID:

EP-2142-M

Laboratory ID: Sampled:

08-29-05 11:30 08-30-05 20:21

Instrument ID: Sample Volume:

MS-6 HP 6890

Received: Extracted: 08-30-05 20:21 09-12-05 14:00 09-21-05 12:28

Final Volume: Percent Solids:

15 g 1 mL

88

1

Analyzed: Analyst:

IIT

Dilution Factor:

CAS Number	Analyte	Concentration Notes	Units	Reporting Limit
91-20-3	Naphthalene	BRL	ug/Kg	11
91-57-6	2-Methylnaphthalene	BRL	ug/Kg	11
208-96-8	Acenaphthylene	BRL	ug/Kg	11
83-32-9	Acenaphthene	BRL	ug/Kg	11
86-73-7	Fluorene	BRL	ug/Kg	11
85-01-8	Phenanthrene	42	ug/Kg	11
120-12-7	Anthracene	BRL	ug/Kg	11
206-44-0	Fluoranthene	110	ug/Kg	11
129-00-0	Pyrene	130	ug/Kg	11
56-55-3	Benzo[a]anthracene	55	ug/Kg	11
218-01-9	Chrysene	63	ug/Kg	11
205-99-2	Benzo[b]fluoranthene	63	ug/Kg	11
207-08-9	Benzo[k]fluoranthene	27	ug/Kg	11
50-32-8	Benzo[a]pyrene	43	ug/Kg	11
193-39-5	Indeno[1,2,3-c,d]pyrene	36	ug/Kg	11
53-70-3	Dibenzo[a,h]anthracene	17	ug/Kg	11
191-24-2	Benzo[g,h,i]perylene	29	ug/Kg	11

QC Surrogate Compound		Measured	Recovery	QC Limits
ortho-Terphenyl	2,900	2,100	72 %	40 - 140 %

Method Reference:

Test Methods for Evaluating Solid Waste, US EPA, SW-846, Third Edition, Update III (1996).

Method modified by use of selected ion monitoring (SIM) in accordance with Section 7.5.5 of the method.

Method protocol modified to include acidification and the surrogate compound in accordance with the MA DEP Method for the Determination of Extractable Petroleum Hydrocarbons. Sample extraction performed by EPA Method 3546. Results are reported on a dry weight basis.

Report Notations:

BRL Indicates concentration, if any, is below reporting limit for analyte. Reporting limit is the lowest concentration that can be reliably quantified under routine laboratory operating conditions. Reporting limits are adjusted for sample size and dilution.



Massachusetts DEP EPH Method Extractable Petroleum Hydrocarbons by GC/FID

Field ID:

W2A02-82905-02

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation: QC Batch ID: Cool

Laboratory ID: Sampled:

87247-02 08-29-05 11:32

Instrument ID: Sample Weight:

EP-2142-M GC-9 Agilent 6890

Received: Extracted:

08-30-05 20:21 09-12-05 14:00

15 g

Analyzed (AL):

09-22-05 06:36

Final Volume:

Analyzed (AR):

09-22-05 07:21

% Solids: Aliphatic Dilution Factor: 1

Analyst:

MM

Aromatic Dilution Factor: 1

EPH Ranges	Concentration Notes	Units	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	36
n-C19 to n-C36 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	36
n-C11 to n-C22 Aromatic Hydrocarbons ^{† 0}	BRL	mg/Kg	36
Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons †	BRL	mg/Kg	36

ractionation:	2-Fluorobiphenyl	3.7	2.2	69 %	40 - 140 %
i lactionation.		3.2	2.2		
	2-Bromonaphthalene	3.2	2.3	<i>7</i> 2 %	40 - 140 %
Extraction:	Chloro-octadecane	3.2	2.2	70 %	40 - 140 %
	ortho -Terphenyl	3.2	2.3	74 %	40 - 140 %

ortho -Terphenyl	3.2	2.3	74 %	40 - 140 %
	650	QA/QC Cer	lification	
Were all QA/QC procedures required by the me Were all performance/acceptance standards for the standards for th			ures achieved?	Yes Yes

Method non-conformances indicated above are detailed below on this data report, or in the accompanying project narrative and project quality control report. Release of this data is authorized by the accompanying signed project cover letter. The accompanying cover letter, project narrative and quality control report are considered part of this data report.

3. Were any significant modifications made to the method, as specified in Section 11.3.1.1?

Method Reference:

Method for the Determination of Extractable Petroleum Hydrocarbons, MA DEP (Revision 1.1, 2004).

Sample extraction performed by microwave accelerated solvent extraction technique. Results are reported on a dry weight basis.

Report Notations:

- Indicates concentration, if any, is below reporting limit for analyte. Reporting limit is the lowest concentration that can be reliably quantified under routine laboratory operating conditions. Reporting limits are adjusted for sample size and dilution.
- Hydrocarbon range data excludes concentrations of any surrogate(s) and/or internal standards eluting in that range.
- n-C11 to n-C22 Aromatic Hydrocarbons range data excludes the method target analyte concentrations.



EPA Method 8270C (Modified) MA DEP EPH Polynuclear Aromatic Hydrocarbons by GC/MS-SIM

Field ID:

W2A02-82905-02

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

GeoInsight, Inc.

Preservation:

Cool

Laboratory ID: Sampled:

: 87247-02 08-29-05 QC Batch ID: Instrument ID: EP-2142-M MS-6 HP 6890

Received:

08-29-05 11:32 08-30-05 20:21 09-12-05 14:00

Sample Volume:

15 g

Extracted: Analyzed:

09-12-05 14:00 09-21-05 13:07 Final Volume: Percent Solids: Dilution Factor:

Analyst: JJT

CAS Number	Analyte	Concentration Notes	Units	Reporting Limit
91-20-3	Naphthalene	BRL	ug/Kg	12
91-57-6	2-Methylnaphthalene	BRL	ug/Kg	12
208-96-8	Acenaphthylene	BRL	ug/Kg	. 12
83-32-9	Acenaphthene	BRL	ug/Kg	12
86-73-7	Fluorene	BRL	ug/Kg	12
85-01-8	Phenanthrene	19	ug/Kg	12
120-12-7	Anthracene	BRL	ug/Kg	12
206-44-0	Fluoranthene	33	ug/Kg	12
129-00-0	Pyrene	29	ug/Kg	12
56-55-3	Benzo[a]anthracene	14	ug/Kg	12
218-01-9	Chrysene	13	ug/Kg	12
205-99-2	Benzo[b]fluoranthene	BRL	ug/Kg	12
207-08-9	Benzo[k]fluoranthene	BRL	ug/Kg	12
50-32-8	Benzo[a]pyrene	BRL	ug/Kg	12
193-39-5	Indeno[1,2,3-c,d]pyrene	12	ug/Kg	12
53-70-3	Dibenzo[a,h]anthracene	BRL	ug/Kg	12
191-24-2	Benzo[g,h,i]perylene	BRL	ug/Kg	12

QC Surrogate Compound	Spiked	Measured	Recovery	QC Limits
ortho-Terphenyl	3,200	2,300	72 %	40 - 140 %

Method Reference:

Test Methods for Evaluating Solid Waste, US EPA, SW-846, Third Edition, Update III (1996).

Method modified by use of selected ion monitoring (SIM) in accordance with Section 7.5.5 of the method.

Method protocol modified to include acidification and the surrogate compound in accordance with the MA DEP Method for the Determination of Extractable Petroleum Hydrocarbons. Sample extraction performed by EPA Method 3546. Results are reported on a dry weight basis.

Report Notations:

BRL Indicates concentration, if any, is below reporting limit for analyte. Reporting limit is the lowest concentration that can be reliably quantified under routine laboratory operating conditions. Reporting limits are adjusted for sample size and dilution.



Project Narrative

Project:

Buzzards Bay/3871-002

Client:

GeoInsight, Inc.

Lab ID:

87247

Received:

08-30-05 20:21

A. Documentation and Client Communication

The following documentation discrepancies, and client changes or amendments were noted for this project:

1 . Samples identified as 'W2A2-82905-01' and 'W2A2-82905-02' were taken off hold and analyzed by MA DEP EPH, per Kevin Trainer, 09-08-05.

B. Method Modifications, Non-Conformances and Observations

The sample(s) in this project were analyzed by the references analytical method(s), and no method modifications, non-conformances or analytical issues were noted, except as indicated below:

1 . MA DEP EPH Note: Samples 87247-01,-02. Polynuclear aromatic hydrocarbon (PAH) target analytes were identified and quantified by GC/MS-SIM, in accordance with the method provision for alternate determinative methodologies. GC/MS-SIM was used to achieve low quantification limits necessary for regulatory compliance. Target analytes were determined utilizing the same sample extract used for carbon range determination by GC/FID.

EUZUUC ON		Herb/PCBs Metals Petraleum Hydrocarbo rable Vol. Ferrirou	AMAZOR SOI	20 T (2) 1 (2) 1 (2) 1 (3) 1 (4) 1 (d.) Thestical in the first in t	PACE IN THE PROPERTY OF THE PR	Inni OHO 3 OHO 3 O	piqipi igsilindi zebporii il 1 kirot il 1 ki	1 SA Herbicides 17 COS/0907 11C	Hell							CHAIN-OF-CUSTODY RECORD	NOTE: All samples submitted subject to Standard Torms and Condition		Time Received by:	Dale Time Received by Laboratory Shipping Airbill Number:	□ GWA Courier □ Express Malt □ Federal Express □ UPS □ Hand □ Number:
CHAIN-OF-CUSTODY REJORD AND WORK ORDER	TURNAROUND	(10 Business Days)	PHIORITY (5 Business Days) PHIORITY (5 Business Days) PHIORITY (6 Business Days)	10) 3	BILLING	33	NPOES	1905	LABONATION (CARACTORIA GOVERN STORE) (CARACT	N	*	* * *	7	7			DATA QUALITY OBJECTIVES	Project Specific QC	Many regulatory programs and EPA methods require project specific OC. Project specific OC includes Sample Duplicates, Matrix Spike Duplicates, Laboratory OC is T. H. M. Project specific or include the project of the project specific includes the project of the project specific includes the project of the project specific includes the project of the	Samples are charged on a per sample basis. Each MS, MSD and Sample Duplicate requires an additional sample aliquot.	Project Specific QC Required Selection of QC Sample Sample Duplicate Chease use sample:	☐ Matrix Spike ☐ Matrix Spike Duplicate ☐ Matrix Spike Duplicate ☐ UPS ☐ Hand
228 Main Street, P.O. Box 1200 Buzzards Bay Ma D2532 Telephone (508) 759-441 • FAX (508) 759-4475 www.goundwateranalytical.com	Firm:	Address:	S Lan Drive	City/State/Zip:	Telephone:	418-649-1114	INSTRUCTIONS: Use separate line for each container (except replicates).	Matrix Type Container(s)	GHOUNDWATER GHOUND	* 1			7					Hegulatory Program	State Standard Deliverables CT XMCP GW-1/S-1 PWS Form	Stand On Stand	☐ Wastewater ☐ Waste Disposal	UVI L'Uredge Material
GROUNDY ER ANALYTICAL	Project Name: Ristande Roll	Project Number:	3871-003	Sampler Name: RW/LC	Project Manager:	Kevin Trainer	INSTRUCTIONS: Use separate line	Sampling	SAMPLE SAMPLE IDENTIFICATION	34-05 1130 WOR 63-839-05-01	50 - 201-00 - 601-00 - 6011	-	-S0402-00408-				MA DEP MCP Data Enhancement Affirmation	VES ON MCB Data Continued	TES IND MCP Drinking Water Sample included. (Require collection of contingent duplicate sample.	Trip blanks are also required, if VOA sample collected). Signature:		



Quality Assurance/Quality Control

A. Program Overview

Groundwater Analytical conducts an active Quality Assurance program to ensure the production of high quality, valid data. This program closely follows the guidance provided by *Interim Guidelines and Specifications for Preparing Quality Assurance Project Plans*, US EPA QAMS-005/80 (1980), and *Test Methods for Evaluating Solid Waste*, US EPA, SW-846, Update III (1996).

Quality Control protocols include written Standard Operating Procedures (SOPs) developed for each analytical method. SOPs are derived from US EPA methodologies and other established references. Standards are prepared from commercially obtained reference materials of certified purity, and documented for traceability.

Quality Assessment protocols for most organic analyses include a minimum of one laboratory control sample, one method blank, one matrix spike sample, and one sample duplicate for each sample preparation batch. All samples, standards, blanks, laboratory control samples, matrix spikes and sample duplicates are spiked with internal standards and surrogate compounds. All instrument sequences begin with an initial calibration verification standard and a blank; and excepting GC/MS sequences, all sequences close with a continuing calibration standard. GC/MS systems are tuned to appropriate ion abundance criteria daily, or for each 12 hour operating period, whichever is more frequent.

Quality Assessment protocols for most inorganic analyses include a minimum of one laboratory control sample, one method blank, one matrix spike sample, and one sample duplicate for each sample preparation batch. Standard curves are derived from one reagent blank and four concentration levels. Curve validity is verified by standard recoveries within plus or minus ten percent of the curve.

B. Definitions

Batches are used as the basic unit for Quality Assessment. A Batch is defined as twenty or fewer samples of the same matrix which are prepared together for the same analysis, using the same lots of reagents and the same techniques or manipulations, all within the same continuum of time, up to but not exceeding 24 hours.

Laboratory Control Samples are used to assess the accuracy of the analytical method. A Laboratory Control Sample consists of reagent water or sodium sulfate spiked with a group of target analytes representative of the method analytes. Accuracy is defined as the degree of agreement of the measured value with the true or expected value. Percent Recoveries for the Laboratory Control Samples are calculated to assess accuracy.

Method Blanks are used to assess the level of contamination present in the analytical system. Method Blanks consist of reagent water or an aliquot of sodium sulfate. Method Blanks are taken through all the appropriate steps of an analytical method. Sample data reported is not corrected for blank contamination.

Surrogate Compounds are used to assess the effectiveness of an analytical method in dealing with each sample matrix. Surrogate Compounds are organic compounds which are similar to the target analytes of interest in chemical behavior, but which are not normally found in environmental samples. Percent Recoveries are calculated for each Surrogate Compound.



Quality Control Report Laboratory Control Samples

LCS

LCSD

Category:

MA DEP EPH Method

Matrix:

QC Batch ID: EP-2142-M

Units:

Soil mg/Kg Instrument ID: Extracted:

Analyzed (AL):

Analyzed (AR):

GC-9 Agilent 6890

09-12-05 14:00

09-13-05 01:09 09-13-05 01:53 Instrument ID: Extracted:

GC-9 Agilent 6890 09-12-05 14:00

Analyzed (AL): Analyzed (AR):

09-13-05 02:37 09-13-05 03:21

Analyst: MM

Analyst:

MM

CAS Number	Analyte		LC	\$		L LC	CS Duplicate	#14.9 - a	QC Lim	its
		Spiked	Measured	Recovery	Spiked	Measured	Recovery	RPD	Spike	RPD
111-84-2	n-Nonane (C ₉)	3.3	1.6	47 %	3.3	1.5	46 %	2 %	30 - 140 %	25%
124-18-5	n-Decane (C ₁₀)	3.3	1.8	55 %	3.3	1.8	54 %	2 %	40 - 140 %	25%
112-40-3	n-Dodecane (C ₁₂)	3.3	1.9	58 %	3.3	1.9	57 %	2 %	40 - 140 %	25%
629-59-4	n-Tetradecane (C ₁₄)	3.3	1.9	59 %	3.3	1.9	59 %	1 %	40 - 140 %	25%
544-76-3	n-Hexadecane (C ₁₆)	3.3	2.2	68 %	3.3	2.2	67 %	2 %	40 - 140 %	25%
593-45-3	n-Octadecane (C ₁₈)	3.3	2.6	80 %	3.3	2.7	81 %	1 %	40 - 140 %	25%
n/a	n-C9 to n-C18 Group	20	12	61 %	20	12	61 %	1 %	40 - 140 %	25%
629-92-5	n-Nonadecane (C ₁₉)	3.3	2.6	80 %	3.3	2.7	82 %	3 %	40 - 140 %	25%
112-95-8	n-Eicosane (C ₂₀)	3.3	2.7	83 %	3.3	2.8	86 %	3 %	40 - 140 %	25%
629-97-0	n-Docosane (C ₂₂)	3.3	2.7	83 %	3.3	2.8	86 %	4 %	40 - 140 %	25%
646-31-1	n-Tetracosane (C ₂₄)	3.3	2.8	84 %	3.3	2.9	87 %	3 %	40 - 140 %	25%
630-01-3	n-Hexacosane (C ₂₆)	3.3	2.7	82 %	3.3	2.8	85 %	3 %	40 - 140 %	25%
630-02-4	n -Octacosane (C ₂₈)	3.3	2.7	82 %	3.3	2.8	85 %	4 %	40 - 140 %	25%
638-68-6	n-Triacontane (C ₃₀)	3.3	2.7	82 %	3.3	2.8	86 %	4 %	40 - 140 %	25%
630-06-8	n-Hexatriacontane (C ₃₆)	3.3	2.4	73 %	3.3	2.5	77 %	5 %	40 - 140 %	25%
n/a	n-C19 to n-C36 Group	26	21	81 %	26	22	84 %	4 %	40 - 140 %	25%
91-20-3	Naphthalene	3.3	2.0	61 %	3.3	1.9	56 %	9 %	40 - 140 %	25%
91-57-6	2-Methylnaphthalene	3.3	2.1	64 %	3.3	1.9	58 %	9 %	40 - 140 %	25%
208-96-8	Acenaphthylene	3.3	2.3	69 %	3.3	2.0	62 %	11 %	40 - 140 %	25%
83-32-9	Acenaphthene	3.3	2.1	65 %	3.3	1.9	58 %	11 %	40 - 140 %	25%
86-73-7	Fluorene	3.3	2.3	71 %	3.3	2.1	63 %	12 %	40 - 140 %	25%
85-01-8	Phenanthrene	3.3	2.7	82 %	3.3	2.3	71 %	14 %	40 - 140 %	25%
120-12-7	Anthracene	3.3	3.0	91 %	3.3	2.8	83 %	8 %	40 - 140 %	25%
206-44-0	Fluoranthene	3.3	3.2	96 %	3.3	2.8	86 %	11 %	40 - 140 %	25%
129-00-0	Pyrene	3.3	3.2	96 %	3.3	2.8	86 %	11 %	40 - 140 %	25%
56-55-3	Benzo[a]anthracene	3.3	3.3	100 %	3.3	3.1	93 %	7 %	40 - 140 %	25%
218-01-9	Chrysene	3.3	3.4	102 %	3.3	3.3	99 %	3 %	40 - 140 %	25%
205-99-2	Benzo[b]fluoranthene	3.3	3.1	94 %	3.3	2.8	86 %	10 %	40 - 140 %	25%
207-08-9	Benzo[k]fluoranthene	3.3	3.2	98 %	3.3	3.1	94 %	4 %	40 - 140 %	25%
50-32-8	Benzo[a]pyrene	3.3	3.1	94 %	3.3	2.9	88 %	7 %	40 - 140 %	25%
193-39-5	Indeno[1,2,3-c,d]pyrene	3.3	2.6	79 %	3.3	2.4	73 %	8 %	40 - 140 %	25%
53-70-3	Dibenzo[a,h]anthracene	3.3	2.8	84 %	3.3	2.7	81 %	4 %	40 - 140 %	25%
191-24-2	Benzo[g,h,i]perylene	3.3	2.4	74 %	3.3	2.3	69 %	6 %	40 - 140 %	25%
n/a	PAH Group	56	47	83 %	56	43	77 %	8 %	40 - 140 %	25%
		57.77 2 228						- 52.00 M 100 C C C C C C C C C C C C C C C C C C		

QC Surrogate	Compound	Spiked	Measured	Recovery	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	2.7	2.1	78 %	2.7	2.0	74 %	40 - 140 %
Ī	2-Bromonaphthalene	2.7	2.2	81 %	2.7	2.1	78 %	40 - 140 %
Extraction:	Chloro-octadecane	2.7	2.1	78 %	2.7	2.1	78 %	40 - 140 %
	ortho-Terphenyl	2.7	2.3	85 %	2.7	1.9	70 %	40 - 140 %

	Frac	tionation Br	eakthrough Evalua	ation -		QC Limits
91-20-3	Naphthalene	LCS	0 %	LCSD	0 %	5%
91-57-6	2-Methylnaphthalene	LCS	0 %	LCSD	0 %	5%

Method Reference:

Method for the Determination of Extractable Petroleum Hydrocarbons, MA DEP (Revision 1.1, 2004).

Method modified by use of microwave accelerated solvent extraction technique.

Report Notations:

All calculations performed prior to rounding. Quality Control Limits are defined by the methodology, or alternatively based upon the historical average recovery plus or minus three standard deviation units.

The LCS and LCSD are prepared from separate source standards than those used for calibration.



Quality Control Report Method Blank

Category: QC Batch ID: MA DEP EPH EP-2142-M

Matrix:

Soil

Instrument ID:

GC-9 Agilent 6890

Extracted:

09-12-05 14:00

Analyzed (AL): Analyzed (AR): 09-12-05 23:40 09-13-05 00:24

Analyst:

MM

EPH Ranges Co	ncentration No	ites Units	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	30
n-C19 to n-C36 Aliphatic Hydrocarbons †	BRL	mg/Kg	30
n-C11 to n-C22 Aromatic Hydrocarbons + 0	BRL	mg/Kg	30
Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons †	BRL	mg/Kg	30

QC Surrogate C	ompound	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	2.7	1.9	72 %	40 - 140 %
	2-Bromonaphthalene	2.7	2.0	74 %	40 - 140 %
Extraction:	Chloro-octadecane	2.7	1.9	73 %	40 - 140 %
	ortho-Terphenyl	2.7	2.0	76 %	40 - 140 %

Method Reference:

Method for the Determination of Extractable Petroleum Hydrocarbons, MA DEP (Revision 1.1, 2004).

Sample extraction performed by microwave accelerated solvent extraction technique.

Report Notations:

BRL Indicates concentration, if any, is below reporting limit for analyte. Reporting limit is the lowest concentration that can be reliably quantified under routine laboratory operating conditions. Reporting limits are adjusted for sample size and dilution.

- † Hydrocarbon range data excludes concentrations of any surrogate(s) and/or internal standards eluting in that range.
- \diamond n-C11 to n-C22 Aromatic Hydrocarbons range data excludes the method target analyte concentrations.



Quality Control Report Laboratory Control Samples

LCSD

Category:

EPA 8270C Modified

Instrument ID:

MS-6 HP 6890

Instrument ID:

MS-6 HP 6890

QC Batch ID: EP-2142-M Matrix:

Soil

Extracted: Analyzed: 09-12-05 14:00 09-13-05 14:03 Extracted: Analyzed: 09-12-05 14:00 09-13-05 14:43

Units:

ug/Kg

Analyst:

JJT

Analyst:

IJΤ

CAS Number	Analyte	5.	LC	S.		-LC	S Duplicate		QC Lim	its
		Spiked	Measured	Recovery	Spiked	Measured	Recovery	RPD	Spike	RPD
91-20-3	Naphthalene	330	220	67 %	330	200	61 %	10 %	40 - 140 %	20%
91-57-6	2-Methylnaphthalene	330	240	73 %	330	220	67 %	9 %	40 - 140 %	20%
85-01-8	Phenanthrene	330	250	76 %	330	230	70 %	8 %	40 - 140 %	20%
83-32-9	Acenaphthene	330	280	85 %	330	250	76 %	11 %	40 - 140 %	20%
208-96-8	Acenaphthylene	330	260	79 %	330	230	70 %	12 %	40 - 140 %	20%
86-73-7	Fluorene	330	260	79 %	330	240	73 %	8 %	40 - 140 %	20%
120-12-7	Anthracene	330	260	79 %	330	240	73 %	8 %	40 - 140 %	20%
206-44-0	Fluoranthene	330	300	91 %	330	290	88 %	3 %	40 - 140 %	20%
129-00-0	Pyrene	330	300	91 %	330	290	88 %	3 %	40 - 140 %	20%
56-55-3	Benzo[a]anthracene	330	310	94 %	330	300	91 %	3 %	40 - 140 %	20%
218-01-9	Chrysene	330	300	91 %	330	290	88 %	3 %	40 - 140 %	20%
205-99-2	Benzo[b]fluoranthene	330	290	88 %	330	280	85 %	4 %	40 - 140 %	20%
207-08-9	Benzo[k]fluoranthene	330	310	94 %	330	290	88 %	7 %	40 - 140 %	20%
50-32-8	Benzo[a]pyrene	330	300	91 %	330	290	88 %	3 %	40 - 140 %	20%
193-39-5	Indeno[1,2,3-c,d]pyrene	330	290	88 %	330	280	85 %	4 %	40 - 140 %	20%
53-70-3	Dibenzo[a,h]anthracene	330	270	82 %	330	260	79 %	4 %	40 - 140 %	20%
191-24-2	Benzo[g,h,i]perylene	330	310	94 %	330	290	88 %	7 %	40 - 140 %	20%
QC Surrogate	Compound	Spiked	Measured	Recovery	Spiked	Measured	Recovery		QC Limi	ts
ortho-Terpher	yl	2,700	2,200	81 %	2,700	2,000	74 %		40 - 140	%

Method Reference:

Test Methods for Evaluating Solid Waste, US EPA, SW-846, Third Edition, Update III (1996).

Method modified by use of selected ion monitoring (SIM) in accordance with Section 7.5.5 of the method.

Method protocol modified to include acidification and the surrogate compound in accordance with the MA DEP Method for the Determination of Extractable Petroleum Hydrocarbons.

Sample extraction performed by EPA Method 3510C.

Report Notations:

All calculations performed prior to rounding. Quality Control Limits are defined by the methodology, or alternatively based upon the historical average recovery plus or minus three standard deviation units.

The LCS and LCSD are prepared from separate source standards than those used for calibration.



Quality Control Report Method Blank

Category:

Matrix:

EPA Method 8270C (Mod.) - EPH PAHs by GC/MS-SIM

QC Batch ID:

EP-2142-M Soil

Instrument ID: Extracted:

MS-6 HP 6890

Analyzed:

09-12-05 14:00 09-13-05 13:23

Analyst:

JJT

CAS Number	Analyte	Concentration Notes	Units	Reporting Limit
91-20-3	Naphthalene	BRL	ug/Kg	10
91-57-6	2-Methylnaphthalene	BRL	ug/Kg	10
208-96-8	Acenaphthylene	BRL	ug/Kg	10
83-32-9	Acenaphthene	BRL	ug/Kg	10
86-73-7	Fluorene	BRL	ug/Kg	10
85-01-8	Phenanthrene	BRL	ug/Kg	10
120-12-7	Anthracene	BRL	ug/Kg	10
206-44-0	Fluoranthene	BRL	ug/Kg	10
129-00-0	Pyrene	BRL	ug/Kg	10
56-55-3	Benzo[a]anthracene	BRL	ug/Kg	10
218-01-9	Chrysene	BRL	ug/Kg	10
205-99-2	Benzo[b]fluoranthene	BRL	ug/Kg	10
207-08-9	Benzo[k]fluoranthene	BRL	ug/Kg	10
50-32-8	Benzo[a]pyrene	BRL	ug/Kg	10
193-39-5	Indeno[1,2,3-c,d]pyrene	BRL	ug/Kg	10
53-70-3	Dibenzo[a,h]anthracene	BRL	ug/Kg	10
191-24-2	Benzo[g,h,i]perylene	BRL	ug/Kg	10

QC Surrogate Compound	Spiked	Measured	Recovery	QC Limits
ortho-Terphenyl	2,700	2,100	78 %	40 - 140 %

Method Reference:

Test Methods for Evaluating Solid Waste, US EPA, SW-846, Third Edition, Update III (1996).

Method modified by use of selected ion monitoring (SIM) in accordance with Section 7.5.5 of the method.

Method protocol modified to include acidification and the surrogate compound in accordance with the MA DEP Method for the Determination of Extractable Petroleum Hydrocarbons.

Sample extraction performed by EPA Method 3546.

Report Notations:

BRL Indicates concentration, if any, is below reporting limit for analyte. Reporting limit is the lowest concentration that can be reliably quantified under routine laboratory operating conditions. Reporting limits are adjusted for sample size and dilution.



Certifications and Approvals

Groundwater Analytical maintains environmental laboratory certification in a variety of states. Copies of our current certificates may be obtained from our website:

http://www.groundwateranalytical.com/qualifications.htm

CONNECTICUT, Department of Health Services, PH-0586

Categories: Potable Water, Wastewater, Solid Waste and Soil

http://www.dph.state.ct.us/BRS/Environmental Lab/OutStateLabList.htm

FLORIDA, Department of Health, Bureau of Laboratories, E87643

Categories: SDWA, CWA, RCRA/CERCLA http://www.floridadep.org/labs/qa/dohforms.htm

MAINE, Department of Human Services, MA103

Categories: Drinking Water and Wastewater

http://www.state.me.us/dhs/eng/water/Compliance.htm

MASSACHUSETTS, Department of Environmental Protection, M-MA-103

Categories: Potable Water and Non-Potable Water http://www.state.ma.us/dep/bspt/wes/files/certlabs.pdf

NEW HAMPSHIRE, Department of Environmental Services, 202703

Categories: Drinking Water and Wastewater

http://www.des.state.nh.us/asp/NHELAP/labsview.asp

NEW YORK, Department of Health, 11754

Categories: Potable Water, Non-Potable Water and Solid Waste

http://www.wadsworth.org/labcert/elap/comm.html

PENNSYLVANIA, Department of Environmental Protection, 68-665

Environmental Laboratory Registration (Non-drinking water and Non-wastewater) http://www.dep.state.pa.us/Labs/Registered/

RHODE ISLAND, Department of Health, 54

Categories: Surface Water, Air, Wastewater, Potable Water, Sewage

http://www.healthri.org/labs/labsCT MA.htm

U.S. Department of Agriculture, Soil Permit, S-53921

Foreign soil import permit

VERMONT, Department of Environmental Conservation, Water Supply Division

Category: Drinking Water

http://www.vermontdrinkingwater.org/wsops/labtable.PDF