GROUNDWATER ANALYTICAL

September 14, 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:

86975

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. Jensen
Operations Manager

EHJ/smd Enclosures



Sample Receipt Report

Project:

Buzzards Bay/3871-002

Client:

Geolnsight, Inc.

Lab ID: 86975

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			vil.	Notes
86975-1	W2A02-P2-M-01		Soil	8/29/05 0:00	MA DEP EPH	with PAHs by 8	8270C-Mod S	IM	
Con 1D	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship		
C668655	120 mL Amber Glass	Industrial	BX17921	None	n/a	n/a	n/a		

Lab ID	Field 1D		Matrix	Sampled	Method				Notes
86975-2	W2A02-P2-M-02		Soil	8/29/05 0:00	MA DEP EPH	with PAHs by	3270C-Mod S	IM	
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship		
C668622	120 mL Amber Glass	Industrial	BX17916	None	n/a	n/a	n/a		

Lab ID	Field 1D		Matrix	Sampled	Method	14.4			Notes
86975-3	W2A02-P2-M-03		Soil	8/29/05 0:00	MA DEP EPH	with PAHs by	8270C-Mod SI	М	
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship		
C668637	120 mL Amber Glass	Industrial	BX17916	None	n/a	n/a	n/a		

Lab ID	Field ID		Matrix	Sampled	Method				Notes
86975-4	W2A02-P2-M-04		Soil	8/29/05 0:00	MA DEP EPH	with PAHs by 8	B270C-Mod S	IM	
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship		
C668614	120 mL Amber Glass	Industrial	BX17916	None	n/a	n/a	n/a		

Lab ID	Field 1D		Matrix	Sampled	Method				Notes	
86975-5	W2A10-P2-M-01		Soil	8/29/05 O:00	MA DEP EPH	with PAHs by	8270C-Mod S	М		
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship			
C668684	120 mL Amber Glass	Industrial	BX17917	None	n/a	n/a	n/a			

Lab ID	Field ID		Matrix	Sampled	Method				en	Notes
86975-6	W2A10-P2-M-02		Soil	8/29/05 0:00	MA DEP EPH	with PAHs b	y 827	OC-Mod S	IM	
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep		Ship		
C668671	120 mL Amber Glass	Industrial	BX17917	None	n/a	n/a		n/a		

Lab ID	Field 1D		Matrix	Sampled	Method				Notes
86975-7	W2A10-P2-M-03		Soil	8/29/05 0:00	MA DEP EPH	with PAHs by	8270C-Mod S	IM	
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship		
C668625	120 mL Amber Glass	Industrial	BX17916	None	n/a	n/a	n/a		

Lab ID	Field ID		Matrix	Sampled	Method			4.45	Notes
86975-8	W2A10-P2-M-04		Soil	8/29/05 0:00	MA DEP EPH	with PAHs by 8	3270C-Mod 5	SIM	
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship		
C668627	120 mL Amber Glass	Industrial	BX17916	None	n/a	n/a	n/a		

Lab 1D	Field 1D		Matrix	Sampled	Method				Notes
86975-9	W2A11-P2-LIT-01		Soil	8/29/05 0:00	MA DEP EPH	with PAHs by 8	3270C-Mod S	IM	
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship		
C668642	120 mL Amber Glass	Industrial	BX17921	None	n/a	n/a	n/a		

Lab ID	Field ID		Matrix	Sampled	Method			Notes
86975-10	W2A11-P2-LIT-02		Soil	8/29/05 0:00	MA DEP EPH	with PAHs by	8270C-Mod SI	IM
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship	
C668647	120 mL Amber Glass	Industrial	BX17921	None	n/a	n/a	n/a	



Sample Receipt Report (Continued)

Client:

Project: Buzzards Bay/3871-002

Geolnsight, Inc.

Lab ID: 86975

Delivery: Hand

Lab Receipt: 08-30-05

Airbill: n/a

Temperature: 4.8'C

Chain of Custody: Present Custody Seal(s): n/a

Lab ID	Field ID		Matrix	Sampled	Method				Notes
86975-11	W2A11-P2-UIT-01		Soil	8/29/05 0:00	MA DEP EPH	with PAHs by i	8270C-Mod S	IM	WEG 25 100
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship		
C668661	120 mL Amber Glass	Industrial	BX17921	None	n/a	n/a	n/a		

Lab ID	Field ID		Matrix	Sampled	Method				Notes
86975-12	W2A11-P2-UIT-02		Soil	8/29/05 0:00	MA DEP EPH	with PAHs by 8	3270C-Mod SI	IM	
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship		
C668644	120 mL Amber Glass	Industrial	BX17921	None	n/a	n/a	n/a		

Lab ID	Field ID		Matrix	Sampled	Method				Notes
86975-13	DDD-P2-01		Soil	8/29/05 0:00	MA DEP EPH	with PAHs by	8270C-Mod SI	М	
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship		
C668685	120 mL Amber Glass	Industrial	BX17917	None	n/a	n/a	n/a		

Lab ID	Field ID	7.7	Matrix	Sampled	Method			Notes
86975-14	DDD-P2-02		Soil	8/29/05 0:00	MA DEP EPH	with PAHs by	8270C-Mod S	SIM
Con ID	Container	Vendor	QC Lot	Preserv	QC Lot	Prep	Ship	
C668674	120 mL Amber Glass	Industrial	BX17917	None	n/a	n/a	n/a	



Field ID:

W2A02-P2-M-01

Project:

Buzzards Bay/3871-002

Client:

Geolnsight, Inc.

Laboratory ID:

86975-01

Sampled: Received: Extracted: 08-29-05 00:00 08-30-05 20:21 09-08-05 11:00

Analyzed (AL): Analyzed (AR):

: 09-12-05 22:20 : 09-12-05 23:04

Analyst:

MM

Matrix:

Soil

Cool

Container:

120 mL Amber Glass

Yes

Yes

Preservation:

QC Batch ID:

EP-2138-M

Instrument ID: Sample Weight:

GC-7 HP 5890 15 g

Final Volume: % Solids:

1 mL 63

Aliphatic Dilution Factor: 1
Aromatic Dilution Factor: 1

EPH Ranges	Concentration Notes	Units -	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons †	BRL	mg/Kg	47
n-C19 to n-C36 Aliphatic Hydrocarbons †	BRL	mg/Kg	47
n-C11 to n-C22 Aromatic Hydrocarbons † 0	52	mg/Kg	47
Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons †	54	mg/Kg	47

QC Surrogate Co	ompound	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	4.2	3.4	82 %	40 - 140 %
	2-Bromonaphthalene	4.2	3.5	84 %	40 - 140 %
Extraction:	Chloro-octadecane	4.2	2.6	61 %	40 - 140 %
	ortho -Terphenyl	4.2	3.9	92 %	40 - 140 %

	A100 (100 CH)	44.00	A STATE OF THE PARTY OF THE PAR	C 2000
(A)	10.00	224 Bar 204 Ba	tificati	120.00
	ALC: NO SECURE	W-98, W-1 G-1 S		223.427

- 1. Were all QA/QC procedures required by the method followed?
- 2. Were all performance/acceptance standards for the required QA/QC procedures achieved?
- 3. Were any significant modifications made to the method, as specified in Section 11.3.1.1?

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.

- 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.
- n-C11 to n-C22 Aromatic Hydrocarbons range data excludes the method target analyte concentrations.



Field ID:

W2A02-P2-M-01

Project:

Buzzards Bay/3871-002

Client: GeoInsight, Inc.

Laboratory ID: Sampled:

86975-01 08-29-05 00:00

Received: Extracted:

08-30-05 20:21 09-08-05 11:00 09-13-05 23:55

Analyzed: Analyst:

IIT

Matrix:

Soil

Container:

120 mL Amber Glass

Preservation:

Cool

QC Batch ID:

EP-2138-M MS-6 HP 6890

Instrument ID: Sample Volume:

15 g

Final Volume:

1 mL

Percent Solids:

63 1

Dilution Factor:

CAS Number	Analyte	Concentration	Notes	Units	Reporting Limit
91-20-3	Naphthalene	15 j	7	ug/Kg	16
91-57-6	2-Methylnaphthalene	BRL		ug/Kg	16
208-96-8	Acenaphthylene	8 j		ug/Kg	16
83-32-9	Acenaphthene	1 5 j		ug/Kg	16
86-73-7	Fluorene	26		ug/Kg	16
85-01-8	Phenanthrene	290	V	ug/Kg	16
120-12-7	Anthracene	36		ug/Kg	16
206-44-0	Fluoranthene	350		ug/Kg	16
129-00-0	Pyrene	350		ug/Kg	16
56-55-3	Benzo[a]anthracene	140		ug/Kg	16
218-01-9	Chrysene	190		ug/Kg	16
205-99-2	Benzo[b]fluoranthene	170		ug/Kg	16
207-08-9	Benzo[k]fluoranthene	64		ug/Kg	16
50-32 - 8	Benzo[a]pyrene	130		ug/Kg	16
193-39-5	Indeno[1,2,3-c,d]pyrene	90		ug/Kg	16
53-70-3	Dibenzo[a,h]anthracene	31		ug/Kg	16
191-24-2	Benzo[g,h,i]perylene	77		ug/Kg	16

QC Surrogate Compound	Spiked	Measured	Recovery	QC Limits
ortho-Terphenyl	4,200	3,500	82 %	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:

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.

Indicates an estimated value detected below the reporting limit for the analyte.



Field ID:

W2A02-P2-M-02

Project:

Buzzards Bay/3871-002

Client:

Geolnsight, Inc.

Laboratory ID:

Geomsigne, me.

86975-02

Sampled: Received: 08-29-05 00:00 08-30-05 20:21 09-08-05 11:00

Extracted: Analyzed (AL): Analyzed (AR):

09-12-05 23:48 09-13-05 00:33

Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons

Analyst:

ММ

Matrix:

Soil

Container:

120 mL Amber Glass

Preservation:

QC Batch ID:

EP-2138-M GC-7 HP 5890

mg/Kg

62

Yes

Yes

Instrument ID: Sample Weight:

15 g 1 mL

Cool

Final Volume: % Solids:

47

Aliphatic Dilution Factor: 1
Aromatic Dilution Factor: 1

EPH Ranges	Concentration No	tes Units	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	62
n-C19 to n-C36 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	62
n-C11 to n-C22 Aromatic Hydrocarbons † 0	BRL	mg/Kg	62

QC Surrogate C	ompound	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	5.5	4.6	82 %	40 - 140 %
	2-Bromonaphthalene	5.5	4.7	84 %	40 - 140 %
Extraction:	Chloro-octadecane	5.5	3.5	64 %	40 - 140 %
	ortho-Terphenyl	5.5	3.6	66 %	40 - 140 %

					Certi		

- 1. Were all QA/QC procedures required by the method followed?
- 2. Were all performance/acceptance standards for the required QA/QC procedures achieved?
- 3. Were any significant modifications made to the method, as specified in Section 11.3.1.1?

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.

BRL

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.

- 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.
- n-C11 to n-C22 Aromatic Hydrocarbons range data excludes the method target analyte concentrations.



Field ID:

W2A02-P2-M-02

Project: Client:

Buzzards Bay/3871-002 GeoInsight, Inc.

Laboratory ID:

86975-02

Sampled: Received: 08-29-05 00:00 08-30-05 20:21

Extracted: Analyzed: Analyst:

09-08-05 11:00 09-14-05 00:34 JJT

Matrix:

Container:

120 mL Amber Glass

Preservation:

Cool

QC Batch ID:

EP-2138-M

Instrument ID:

MS-6 HP 6890

Sample Volume:

15 g 1 mL

Final Volume: Percent Solids:

47 Dilution Factor:

CAS Number	Analyte	Concentration	Notes	Units	Reporting Limit
91-20-3	Naphthalene	46		ug/Kg	21
91-57-6	2-Methylnaphthalene	24		ug/Kg	21
208-96-8	Acenaphthylene	35		ug/Kg	21
83-32-9	Acenaphthene	48		ug/Kg	21
86-73-7	Fluorene	58		ug/Kg	21
85-01-8	Phenanthrene	630		ug/Kg	21
120-12-7	Anthracene	130		ug/Kg	21
206-44-0	Fluoranthene	1,000		ug/Kg	21
129-00-0	Pyrene	980		ug/Kg	21
56-55-3	Benzo[a]anthracene	470		ug/Kg	21
218-01-9	Chrysene	510		ug/Kg	21
205-99-2	Benzo[b]fluoranthene	550		ug/Kg	21
207-08-9	Benzo[k]fluoranthene	210		ug/Kg	21
50-32-8	Benzo[a]pyrene	540		ug/Kg	21
193-39-5	Indeno[1,2,3-c,d]pyrene	370		ug/Kg	21
53-70-3	Dibenzo[a,h]anthracene	82		ug/Kg	21
191-24-2	Benzo[g,h,i]perylene	380		ug/Kg	21

QC Surrogate Compound	Spiked	Measured	Recovery	QCLimits
ortho- Terphenyl	5,500	3,800	69 %	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:

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.



Field ID:

W2A02-P2-M-03

Project:

Buzzards Bay/3871-002

Client:

Received:

Geolnsight, Inc.

Laboratory ID: Sampled:

86975-03 08-29-05 00:00 08-30-05 20:21 09-08-05 11:00

Extracted: Analyzed (AL):

09-13-05 01:17 Analyzed (AR): 09-13-05 02:01

Analyst:

MM

Soil

Container:

120 mL Amber Glass

Preservation: Cool

QC Batch ID:

EP-2138-M GC-7 HP 5890

Instrument ID: Sample Weight:

16 g

Final Volume: % Solids:

1 mL 82

Aliphatic Dilution Factor: 1 Aromatic Dilution Factor: 1

EPH Ranges C	oncentration No	tes Units	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	35
n-C19 to n-C36 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	35
n-C11 to n-C22 Aromatic Hydrocarbons † 0	BRL	mg/Kg	35
Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons †	BRL	mg/Kg	35

QC Surrogate C	ompound	Spiked	Measured	Recovery	OC Limits
Fractionation:	2-Fluorobiphenyl	3.2	2.4	75 %	40 - 140 %
	2-Bromonaphthalene	3.2	2.4	76 %	40 - 140 %
Extraction:	Chloro-octadecane	3.2	1.7	53 %	40 - 140 %
	ortho-Terphenyl	3.2	1.7	53 %	40 - 140 %

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?

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

Yes 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.

- 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.



Field ID:

W2A02-P2-M-03

Matrix:

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID:

86975-03

QC Batch ID:

EP-2138-M

Sampled: Received: 08-29-05 00:00 08-30-05 20:21

Instrument ID: Sample Volume: MS-6 HP 6890

Extracted:

09-08-05 11:00

16 g

Analyzed:

09-14-05 01:13

Final Volume: Percent Solids: 1 mL 82

Analyst:

JJT

Dilution Factor:

1

CAS Number	Analyte	Concentration Notes	Units	Reporting Limit
91-20-3	Naphthalene	14	ug/Kg	12
91-57-6	2-Methylnaphthalene	7 j	ug/Kg	12
208-96-8	Acenaphthylene	14	ug/Kg	12
83-32-9	Acenaphthene	13	ug/Kg	12
86-73-7	Fluorene	24	ug/Kg	12
85-01-8	Phenanthrene	260	ug/Kg	12
120-12-7	Anthracene	56	ug/Kg	12
206-44-0	Fluoranthene	400	ug/Kg	12
129-00-0	Pyrene	430	ug/Kg	12
56-55-3	Benzo[a]anthracene	220	ug/Kg	12
218-01-9	Chrysene	220	ug/Kg	12
205-99-2	Benzo[b]fluoranthene	200	ug/Kg	12
207-08-9	Benzo[k]fluoranthene	72	ug/Kg	12
50-32-8	Benzo[a]pyrene	200	ug/Kg	12
193-39-5	Indeno[1,2,3-c,d]pyrene	120	ug/Kg	12
53-70-3	Dibenzo[a,h]anthracene	35	ug/Kg	12
191-24-2	Benzo[g,h,i]perylene	120	ug/Kg	12

QC Surrogate Compound	Spiked	Measured	Recovery	QC Limits
ortho-Terphenyl	3,200	1,800	57 %	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:

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.

Indicates an estimated value detected below the reporting limit for the analyte.



Field ID:

W2A02-P2-M-04

Project:

Buzzards Bay/3871-002

Client:

Geolnsight, Inc.

Laboratory ID:

86975-04

Sampled: Received: 08-29-05 00:00 08-30-05 20:21 09-08-05 11:00

Extracted: Analyzed (AL):

09-13-05 02:45 09-13-05 03:29 Analyzed (AR):

Analyst:

MM

Matrix.

Container:

120 mL Amber Glass

Preservation: Cool

QC Batch ID:

Instrument ID:

GC-7 HP 5890

Sample Weight: Final Volume:

16 g 1 mL 82

EP-2138-M

Soil

% Solids: Aliphatic Dilution Factor: 1

Aromatic Dilution Factor: 1

EPH Ranges Co	oncentration No	ites Units	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons †	BRL	mg/Kg	35
n-C19 to n-C36 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	35
n-C11 to n-C22 Aromatic Hydrocarbons + 0	BRL	mg/Kg	35
Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons †	BRL	mg/Kg	35

QC Surrogate Co	ompound	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	3.1	2.5	80 %	40 - 140 %
	2-Bromonaphthalene	3.1	2.7	87 %	40 - 140 %
Extraction:	Chloro-octadecane	3.1	2.3	73 %	40 - 140 %
	ortho-Terphenyl	3.1	2.5	80 %	40 - 140 %

- 1. Were all QA/QC procedures required by the method followed?
- 2. Were all performance/acceptance standards for the required QA/QC procedures achieved? 3. Were any significant modifications made to the method, as specified in Section 11.3.1.1?

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.

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.

- 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.
- ٥ n-C11 to n-C22 Aromatic Hydrocarbons range data excludes the method target analyte concentrations.



Field ID:

W2A02-P2-M-04

Matrix:

Cool

82

Project:

Buzzards Bay/3871-002

Container: Preservation: 120 mL Amber Glass

Client:

GeoInsight, Inc.

Laboratory ID:

86975-04

QC Batch ID:

EP-2138-M

Sampled:

08-29-05 00:00

Instrument ID:

MS-6 HP 6890

Received:

08-30-05 20:21

Extracted:

09-08-05 11:00

Sample Volume:

Analyzed:

09-14-05 01:52

Final Volume: Percent Solids: 16 g 1 mL

Analyst:

JJT

Dilution Factor:

CAS Number	Analyte	Concentration Notes	Units	Reporting Limit
91-20-3	Naphthalene	7 . j	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	10 j	ug/Kg	12
120-12-7	Anthracene	BRL	ug/Kg	12
206-44-0	Fluoranthene	28	ug/Kg	12
129-00-0	Pyrene	29	ug/Kg	12
56-55-3	Benzo[a]anthracene	12	ug/Kg	12
218-01-9	Chrysene	12	ug/Kg	12
205-99-2	Benzo[b]fluoranthene	6 j	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	13	ug/Kg	12
53-70-3	Dibenzo[a,h]anthracene	13	ug/Kg	12
191-24-2	Benzo[g,h,i]perylene	8 j	ug/Kg	12

QC Surrogate Compound	Spiked	Measured	Recovery	QC Limits
ortho-Terphenyl	3,100	2,400	77 %	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:

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.

Indicates an estimated value detected below the reporting limit for the analyte.



Field ID:

W2A10-P2-M-01

Project:

Buzzards Bay/3871-002

Client:

Geolnsight, Inc.

86975-05

Laboratory ID: Sampled: Received:

08-29-05 00:00 08-30-05 20:21 09-08-05 11:00

Extracted: Analyzed (AL):

09-12-05 14:09 09-12-05 14:49 Analyzed (AR):

Analyst:

MM

Matrix:

Cool

Container:

120 mL Amber Glass

Preservation:

OC Batch ID:

Instrument ID:

Sample Weight: Final Volume:

16 g 1 mL

% Solids:

59

EP-2138-M

GC-7 HP 5890

Aliphatic Dilution Factor: 1 Aromatic Dilution Factor: 1

EPH Ranges	Concentration Notes	Units	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	49
n-C19 to n-C36 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	49
n-C11 to n-C22 Aromatic Hydrocarbons + 0	62	mg/Kg	49
Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons †	63	mg/Kg	49

QC Surrogate Co	ompound	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	4.4	3.5	80 %	40 - 140 %
	2-Bromonaphthalene	4.4	3.8	86 %	40 - 140 %
Extraction: Chloro-oc	Chloro-octadecane	4.4	3.1	71 %	40 - 140 %
	ortho-Terphenyl	4.4	4.4	100 %	40 - 140 %

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?

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.

- 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. 0



Field ID:

W2A10-P2-M-01

Matrix:

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID:

86975-05

QC Batch ID:

EP-2138-M

Sampled:

08-29-05 00:00

Instrument ID:

MS-6 HP 6890

Received:

08-30-05 20:21

Sample Volume:

Extracted:

09-08-05 11:00 09-14-05 02:32

Final Volume: Percent Solids: 16 g 1 mL

59

1

Analyzed: Analyst:

JJT

Dilution Factor:

CAS Number	Analyte	Concentration Notes	Units	Reporting Limit
91-20-3	Naphthalene	10 j	ug/Kg	16
91-5 <i>7</i> -6	2-Methylnaphthalene	BRL	ug/Kg	16
208-96-8	Acenaphthylene	BRL	ug/Kg	16
83-32-9	Acenaphthene	BRL	ug/Kg	16
86-73-7	Fluorene	BRL	ug/Kg	16
85-01-8	Phenanthrene	12 j	ug/Kg	16
120-12-7	Anthracene	BRL	ug/Kg	16
206-44-0	Fluoranthene	31	ug/Kg	16
129-00-0	Pyrene	33	ug/Kg	16
56-55-3	Benzo[a]anthracene	16	ug/Kg	16
218-01-9	Chrysene	90	ug/Kg	16
205-99-2	Benzo[b]fluoranthene	49	ug/Kg	16
207-08-9	Benzo[k]fluoranthene	9 j	ug/Kg	16
50-32-8	Benzo[a]pyrene	66	ug/Kg	16
193-39-5	Indeno[1,2,3-c,d]pyrene	34	ug/Kg	16
53-70-3	Dibenzo[a,h]anthracene	31	ug/Kg	16
191-24-2	Benzo[g,h,i]perylene	42	ug/Kg	16

QC Surrogate Compound	Spiked	Measured	Recovery	QC Limits
ortho-Terphenyl	4,400	3,200	73 %	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:

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.

Indicates an estimated value detected below the reporting limit for the analyte.



Field ID:

W2A10-P2-M-02

Project:

Buzzards Bay/3871-002

Client:

Geolnsight, Inc.

Laboratory ID: Sampled:

08-29-05 00:00 08-30-05 20:21

86975-06

Received: Extracted: Analyzed (AL):

09-08-05 11:00 09-12-05 15:33 09-12-05 16:18

Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons

Analyzed (AR): Analyst:

MM

Matrix: Container: Soil

120 mL Amber Glass

Preservation:

Cool

QC Batch ID:

Instrument ID:

EP-2138-M GC-7 HP 5890

mg/Kg

Sample Weight:

15 g

Final Volume: % Solids:

1 mL 77

Aliphatic Dilution Factor: 1 Aromatic Dilution Factor: 1

EPH Ranges	Concentration Notes	Units	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	38
n-C19 to n-C36 Aliphatic Hydrocarbons †	BRL	mg/Kg	38
n-C11 to n-C22 Aromatic Hydrocarbons † 0	BRL	mg/Kg	38

QC Surrogate C	ompound	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	3.4	2.6	78 %	40 - 140 %
	2-Bromonaphthalene	3.4	2.8	82 %	40 - 140 %
Extraction:	Chloro-octadecane	3.4	2.7	80 %	40 - 140 %
	ortho -Terphenyl	3.4	2.8	82 %	40 - 140 %

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

- Yes No

Yes

38

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.

- 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. ٥



Field ID:

2A10-P2-M-02

Matrix:

Soil

Project: Client: Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Geolnsight, Inc. 86975-06

Preservation: QC Batch ID: Cool

Laboratory ID: Sampled:

08-29-05 00:00

Instrument ID:

EP-2138-M MS-6 HP 6890

Received:

08-30-05 20:21 09-08-05 11:00

Sample Volume:

15 g

Extracted: Analyzed:

09-14-05 03:11

Final Volume: Percent Solids:

1 mL 77

Analyst:

JJT

Dilution Factor:

ctor: 1

CAS Number	Analyte	Concentration Notes	Units	Reporting Limit
91-20-3	Naphthalene	BRL	ug/Kg	18
91-57-6	2-Methylnaphthalene	BRL	ug/Kg	18
208-96-8	Acenaphthylene	BRL	ug/Kg	18
83-32-9	Acenaphthene	BRL	ug/Kg	18
86-73-7	Fluorene	BRL	ug/Kg	18
85-01-8	Phenanthrene	BRL	ug/Kg	18
120-12-7	Anthracene	BRL	ug/Kg	18
206-44-0	Fluoranthene	BRL	ug/Kg	18
129-00-0	Pyrene	BRL	ug/Kg	18
56-55-3	Benzo[a]anthracene	BRL	ug/Kg	18
218-01-9	Chrysene	BRL	ug/Kg	18
205-99-2	Benzo[b]fluoranthene	BRL	ug/Kg	18
207-08-9	Benzo[k]fluoranthene	BRL	ug/Kg	18
50-32-8	Benzo[a]pyrene	BRL	ug/Kg	18
193-39-5	Indeno[1,2,3-c,d]pyrene	BRL	ug/Kg	18
53-70-3	Dibenzo[a,h]anthracene	BRL	ug/Kg	18
191-24-2	Benzo[g,h,i]perylene	BRL	ug/Kg	18

QC Surrogate Compound	Spiked	Measured	Recovery	QC Limits
ortho-Terphenyl	3,400	2,700	81 %	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:

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.



Field ID:

W2A10-P2-M-03

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID: Sampled:

08-29-05 00:00

86975-07

QC Batch ID: Instrument ID: EP-2138-M

mg/Kg

Received:

08-30-05 20:21 09-08-05 11:00

Sample Weight:

GC-7 HP 5890 15 g

Extracted: Analyzed (AL):

09-12-05 17:02

Final Volume:

1 mL

Analyzed (AR):

09-12-05 17:47

Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons

% Solids: Aliphatic Dilution Factor: 1

Analyst:

MM

Aromatic Dilution Factor: 1

BRL

EPH Ranges	Concentration No	tes Units	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	56
n-C19 to n-C36 Aliphatic Hydrocarbons †	BRL	mg/Kg	56
n-C11 to n-C22 Aromatic Hydrocarbons † °	BRL	mg/Kg	56

QC Surrogate Co	ompound	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	5.0	4.0	80 %	40 - 140 %
	2-Bromonaphthalene	5.0	4.0	80 %	40 - 140 %
Extraction:	Chloro-octadecane	5.0	3.7	75 %	40 - 140 %
	ortho-Terphenyl	5.0	4.5	90 %	40 - 140 %

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

56

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.

- 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.



Field ID:

W2A10-P2-M-03

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID:

86975-07

QC Batch ID: Instrument ID: EP-2138-M

Sampled: Received: 08-29-05 00:00 08-30-05 20:21

Sample Volume:

MS-6 HP 6890

Extracted:

09-08-05 11:00

15 g

Analyzed:

09-14-05 03:50

Final Volume:

1 mL

Analyst:

JJT

Percent Solids:

52 Dilution Factor: 1

CAS Number	Analyte	Concentration	Notes Units	Reporting Limit
91-20-3	Naphthalene	10 j	ug/Kg	19
91-57-6	2-Methylnaphthalene	BRL	ug/Kg	19
208-96-8	Acenaphthylene	BRL	ug/Kg	19
83-32-9	Acenaphthene	BRL	ug/Kg	19
86-73-7	Fluorene	BRL	ug/Kg	19
85-01-8	Phenanthrene	BRL	ug/Kg	19
120-12-7	Anthracene	BRL	ug/Kg	19
206-44-0	Fluoranthene	15 j	ug/Kg	19
129-00-0	Pyrene	17 j	ug/Kg	19
56-55-3	Benzo[a]anthracene	BRL	ug/Kg	19
218-01-9	Chrysene	BRL	ug/Kg	19
205-99-2	Benzo[b]fluoranthene	BRL	ug/Kg	19
207-08-9	Benzo[k]fluoranthene	BRL	ug/Kg	19
50-32-8	Benzo[a]pyrene	BRL	ug/Kg	19
193-39-5	Indeno[1,2,3-c,d]pyrene	16 j	ug/Kg	19
53-70-3	Dibenzo[a,h]anthracene	21	ug/Kg	19
191-24-2	Benzo[g,h,i]perylene	BRL	ug/Kg	19

QC Surrogate Compound	Spiked	Measured	Recovery	QC:Limits
ortho-Terphenyl	5,000	4,000	81 %	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:

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.

Indicates an estimated value detected below the reporting limit for the analyte.



Field ID:

W2A10-P2-M-04

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID: Sampled:

86975-08 08-29-05 00:00 OC Batch ID: Instrument ID: EP-2138-M

Received: Extracted: 08-30-05 20:21 09-08-05 11:00

Sample Weight:

GC-7 HP 5890 16 g

Analyzed (AL):

09-12-05 18:31

Final Volume:

1 mL 77

Analyzed (AR): Analyst:

09-12-05 19:16

мм

% Solids:

Aliphatic Dilution Factor: Aromatic Dilution Factor: 1

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	the state of the s
EPH Rang	es

EPH Ranges (Concentration Note	es Units	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	38
n-C19 to n-C36 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	38
n-C11 to n-C22 Aromatic Hydrocarbons ^{† 0}	BRL	mg/Kg	38
Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons †	BRL	mg/Kg	38

<u>Unadjusted</u> n-C11 to n-C22 Aromatic Hydrocarbons [†]				BRL	mg/Kg	mg/Kg 38	
QC Surrogate G	ompound	Spiked	Measured	Recovery	Q	Limits	
Fractionation:	2-Fluorobiphenyl	3.3	2.6	79 %	40	- 140 %	
	2-Bromonaphthalene	3.3	2.8	85 %	40	- 140 %	
Extraction:	Chloro-octadecane	3.3	2.7	80 %	40	- 140 %	
	ortho-Terphenyl	3.3	2.7	80 %	40	- 140 %	

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

Yes Yes 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.

- 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.
- n-C11 to n-C22 Aromatic Hydrocarbons range data excludes the method target analyte concentrations.



Field ID:

W2A10-P2-M-04

Matrix:

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation: QC Batch ID:

Cool EP-2138-M

Laboratory ID: Sampled:

86975-08 08-29-05 00:00

Instrument ID:

MS-6 HP 6890

Received:

08-30-05 20:21

Sample Volume:

16 g

Extracted: Analyzed: 09-08-05 11:00 09-14-05 04:29 Final Volume:

1 mL 77

Analyst:

JJT

Percent Solids: Dilution Factor:

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

QC Surrogate Compound	Spiked	Measured	1Recovery	QC Limits
ortho- Terphenyl	3,300	2,500	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 3546. 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.



Field ID:

W2A11-P2-LIT-01

Project:

Buzzards Bay/3871-002

Client:

GeoInsight, Inc.

Laboratory ID:

86975-09

Sampled: Received: 08-29-05 00:00 08-30-05 20:21 09-08-05 11:00

Extracted: Analyzed (AL):

): 09-12-05 20:00): 09-12-05 20:45

Analyzed (AR): Analyst:

MM

Matrix:

Soil

Container:

120 mL Amber Glass

Preservation:

Cool

QC Batch ID:

Instrument ID:

EP-2138-M GC-7 HP 5890

Sample Weight:

15 g 1 mL

Final Volume:

ide.

% Solids: 83
Aliphatic Dilution Factor: 1

Aromatic Dilution Factor: 1

EPH Ranges	oncentration Note	es 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

QC Surrogate G	ompound	Spiked	Measured	Recovery	OC Limits
Fractionation:	2-Fluorobiphenyl	3.2	2.5	79 %	40 - 140 %
	2-Bromonaphthalene	3.2	2.6	83 %	40 - 140 %
Extraction:	Chloro-octadecane	3.2	2.4	76 %	40 - 140 %
	ortho-Terphenyl	3.2	2.4	76 %	40 - 140 %

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

No

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

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.

- 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.
- n-C11 to n-C22 Aromatic Hydrocarbons range data excludes the method target analyte concentrations.



Field ID:

W2A11-P2-LIT-01

Matrix:

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID:

86975-09

QC Batch ID:

EP-2138-M

Sampled: Received: 08-29-05 00:00 08-30-05 20:21

Instrument ID: Sample Volume: MS-6 HP 6890 15 g

Extracted: Analyzed: 09-08-05 11:00 09-14-05 05:08 Final Volume:

1 mL 83

1

Analyst:

JJT

Percent Solids:

Dilution Factor:

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	BRL	ug/Kg	12
120-12-7	Anthracene	BRL	ug/Kg	12
206-44-0	Fluoranthene	BRL	ug/Kg	12
129-00-0	Pyrene	BRL	ug/Kg	12
56-55-3	Benzo[a]anthracene	BRL	ug/Kg	12
218-01-9	Chrysene	BRL	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	BRL	ug/Kg	12
53- <i>7</i> 0-3	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	OC Limits
ortho-Terphenyl	3,200	2,400	77 %	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.



Field ID:

W2A11-P2-LIT-02

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID: Sampled:

86975-10

OC Batch ID:

EP-2138-M

Received:

08-29-05 00:00

Instrument ID: Sample Weight: GC-7 HP 5890

Extracted:

08-30-05 20:21 09-08-05 11:00

Final Volume:

15 g

Analyzed (AL):

09-12-05 22:13

% Solids:

1 mL

Analyzed (AR):

09-12-05 22:57

Aliphatic Dilution Factor: 1

Analyst:

MM

Aromatic Dilution Factor: 1

EPH Ranges - Go	oncentration No	tes 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 Co	ompound	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	2.7	2.5	93 %	40 - 140 %
	2-Bromonaphthalene	2.7	2.6	96 %	40 - 140 %
Extraction:	Chloro-octadecane	2.7	2.1	79 %	40 - 140 %
	ortho-Terphenyl	2.7	2.3	86 %	40 - 140 %

Extraction:	Chioro-octadecane	2.7	2.1	7 9 %	40 - 140 %
	ortho -Terphenyl	2.7	2.3	86 %	40 - 140 %
	the color		QA/QC Cert	ification	
1. Were all QA/	QC procedures required by the m	nethod followed	d?		Yes

2. Were all performance/acceptance standards for the required QA/QC procedures achieved? 3. Were any significant modifications made to the method, as specified in Section 11.3.1.1?

Yes 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.

- 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.



Field ID:

W2A11-P2-LIT-02

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID:

86975-10

QC Batch ID:

EP-2138-M

Sampled: Received:

08-29-05 00:00 08-30-05 20:21

Instrument ID: Sample Volume: MS-6 HP 6890 15 g

Extracted:

09-08-05 11:00 09-14-05 05:47

Final Volume:

1 mL 97

1

Analyzed: Analyst:

JJT

Percent Solids:

Dilution Factor:

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	2,300	85 %	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:

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.



Field ID: Project:

W2A11-P2-UIT-01

Buzzards Bay/3871-002

Client:

GeoInsight, Inc.

86975-11

Laboratory ID: Sampled: Received:

08-29-05 00:00 08-30-05 20:21

Extracted: Analyzed (AL):

09-09-05 15:00 09-12-05 23:42 09-13-05 00:26

Analyzed (AR): Analyst:

MM

Matrix:

Soil

Container:

120 mL Amber Glass Cool

Preservation:

QC Batch ID:

EP-2140-M GC-7 HP 5890

Instrument ID:

Sample Weight: 16 g

Final Volume:

1 mL

% Solids:

95

Aliphatic Dilution Factor: 1 Aromatic Dilution Factor: 1

EPH Ranges	Concentration Note	s 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	OC Limits
Fractionation:	2-Fluorobiphenyl	2.7	2.0	75 %	40 - 140 %
	2-Bromonaphthalene	2.7	2.1	79 %	40 - 140 %
Extraction:	Chloro-octadecane	2.7	2.0	76 %	40 - 140 %
	ortho -Terphenyl	2.7	1.9	71 %	40 - 140 %

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

Yes Yes

Νo

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.

- 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.



Field ID:

W2A11-P2-UIT-01

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID: Sampled:

86975-11

QC Batch ID: Instrument ID: EP-2140-M

Received:

08-29-05 00:00 08-30-05 20:21

Sample Volume:

MS-6 HP 6890

Extracted: Analyzed: 09-09-05 15:00 09-14-05 06:26

Final Volume:

16 g 1 mL

Analyst:

Percent Solids: Dilution Factor:

95 1

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	6 j	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	7 j	ug/Kg	10
53-70-3	Dibenzo[a,h]anthracene	11	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	1,900	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:

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.

Indicates an estimated value detected below the reporting limit for the analyte.



Field ID:

W2A11-P2-UIT-02

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container: Preservation:

120 mL Amber Glass Cool

Client:

GeoInsight, Inc.

86975-12

QC Batch ID:

EP-2140-M

Laboratory ID: Sampled: Received:

08-29-05 00:00 08-30-05 20:21

Instrument ID: Sample Weight: GC-7 HP 5890

Extracted: Analyzed (AL): 09-09-05 15:00 09-13-05 01:10 Final Volume:

15 g 1 mL 94

mg/Kg

Analyzed (AR):

09-13-05 01:54

Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons 1

% Solids:

BRL

Analyst:

MM

Aliphatic Dilution Factor: 1 Aromatic Dilution Factor: 1

EPH Ranges	Concentration Notes	Units	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	31
n-C19 to n-C36 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	31
n-C11 to n-C22 Aromatic Hydrocarbons † 0	BRL	mg/Kg	31

QC Surrogate C	ompound	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	2.8	2.4	88 %	40 - 140 %
	2-Bromonaphthalene	2.8	2.6	95 %	40 - 140 %
Extraction:	Chloro-octadecane	2.8	2.1	75 %	40 - 140 %
	ortho -Terphenyl	2.8	2.3	84 %	40 - 140 %

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 No

31

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

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.

- 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.



Field ID:

W2A11-P2-UIT-02

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID: Sampled:

86975-12

QC Batch ID:

EP-2140-M

Received:

08-29-05 00:00

Instrument ID: Sample Volume: MS-6 HP 6890

Extracted:

08-30-05 20:21 09-09-05 15:00

Final Volume:

15 g 1 mL

Analyzed:

09-14-05 07:05

Percent Solids: Dilution Factor: 94

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	BRL	ug/Kg	12
120-12-7	Anthracene	BRL	ug/Kg	12
206-44-0	Fluoranthene	BRL	ug/Kg	12
129-00-0	Pyrene	BRL	ug/Kg	12
56-55-3	Benzo[a]anthracene	BRL	ug/Kg	12
218-01-9	Chrysene	BRL	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	BRL	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	2,800	2,200	79 %	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:

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.



Field ID: Project:

DDD-P2-01

Matrix:

Soil

Cool

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

EP-2140-M

Laboratory ID: Sampled:

86975-13 08-29-05 00:00 QC Batch ID: Instrument ID:

Received:

08-30-05 20:21

Sample Weight:

GC-7 HP 5890

Extracted:

09-09-05 15:00 09-13-05 02:38 Final Volume:

16 g 1 mL

mg/Kg

Analyzed (AL): Analyzed (AR):

09-13-05 03:22

Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons †

% Solids:

86

Analyst:

MM

Aliphatic Dilution Factor: 1 Aromatic Dilution Factor: 1

BRL

EPH Ranges	Concentration Not	es 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 † 0	BRL	mg/Kg	33

QC Surrogate G	ompound	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	3.0	2.4	80 %	40 - 140 %
	2-Bromonaphthalene	3.0	2.5	84 %	40 - 140 %
Extraction:	Chloro-octadecane	3.0	2.3	77 %	40 - 140 %
	ortho-Terphenyl	3.0	2.3	76 %	40 - 140 %

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 No

33

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

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.

- 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.
- ٥ n-C11 to n-C22 Aromatic Hydrocarbons range data excludes the method target analyte concentrations.



Field ID:

DDD-P2-01

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID:

86975-13

QC Batch ID:

EP-2140-M

Sampled:

08-29-05 00:00

Instrument ID:

MS-6 HP 6890

Received:

08-30-05 20:21

Sample Volume:

Extracted:

09-09-05 15:00

Final Volume:

16 g 1 mL

Analyzed:

09-14-05 07:44

Percent Solids: Dilution Factor: 86 1

Analyst:

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

QC Surrogate Compound	Spiked	Measured	Recovery	100	QC Limits
ortho-Terphenyl	3,000	2,400	80 %		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:

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.



Field ID:

DDD-P2-02

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID: Sampled:

86975-14 08-29-05 00:00

OC Batch ID: Instrument ID: EP-2140-M GC-7 HP 5890

Received:

08-30-05 20:21

Sample Weight:

Extracted:

09-09-05 15:00

Final Volume:

15 g

mg/Kg

Analyzed (AL):

09-13-05 04:07

1 ml 93

Analyzed (AR):

09-13-05 04:51

Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons †

% Solids:

Aliphatic Dilution Factor: 1

BRL

Analyst:

MM

Aromatic Dilution Factor: 1

EPH Ranges	Concentration Note	es Units	Reporting Limit
n-C9 to n-C18 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	32
n-C19 to n-C36 Aliphatic Hydrocarbons [†]	BRL	mg/Kg	32
n-C11 to n-C22 Aromatic Hydrocarbons † °	BRL	mg/Kg	32

QC Surrogate C	ompound	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	2.8	2.3	83 %	40 - 140 %
	2-Bromonaphthalene	2.8	2.2	77 %	40 - 140 %
Extraction:	Chloro-octadecane	2.8	2.0	69 %	40 - 140 %
	ortho-Terphenyl	2.8	2.3	81 %	40 - 140 %

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

32

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.

- 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.



Field ID:

DDD-P2-02

Matrix:

Soil

Project:

Buzzards Bay/3871-002

Container:

120 mL Amber Glass

Client:

Geolnsight, Inc.

Preservation:

Cool

Laboratory ID:

86975-14

QC Batch ID: Instrument ID:

EP-2140-M

Sampled: Received: 08-29-05 00:00 08-30-05 20:21

Sample Volume:

MS-6 HP 6890 15 g

Extracted: Analyzed:

09-09-05 15:00 09-14-05 08:23

Benzo[b]fluoranthene

Benzo[k]fluoranthene

Indeno[1,2,3-c,d]pyrene

Dibenzo[a,h]anthracene

Benzo[a]pyrene

BRL

Final Volume:

1 mL 93

ug/Kg

ug/Kg

ug/Kg

ug/Kg

ug/Kg

12

12

12

12

12

Analyst:

JJT

Percent Solids: Dilution Factor:

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	BRL	ug/Kg	12
120-12-7	Anthracene	BRL	ug/Kg	12
206-44-0	Fluoranthene	BRL	ug/Kg	12
129-00-0	Pyrene	BRL	ug/Kg	12
56-55-3	Benzo[a]anthracene	BRL	ug/Kg	12
218-01-9	Chrysene	BRL	ug/Kg	12

191-24-2	Benzo[g,h,i]perylene			BRL	ug/Kg	12
QC Surrogate C	ompound	Spiked	Measured	Recovery	Q	C Limits
ortho-Terpheny		2,800	2,300	83 %	40	- 140 %

Method Reference:

205-99-2

207-08-9

50-32-8

193-39-5

53-70-3

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:

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.

BRL

BRL

BRL

BRL

BRL



Project Narrative

Project:

Buzzards Bay/3871-002

Client:

Geolnsight, Inc.

Lab ID:

86975

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. No documentation discrepancies, changes, or amendments were noted.

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. No method modifications, non-conformances or analytical issues were noted.

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GROUNDWATER ANALYTICAL	228 Main Street, P.O. Box 1200 Buzzards Bay, MA 02532 Telephone (508) 759-4441 • FAX (508) 759-4475	CHAIN-OF-CUSTODY RECORD AND WORK ORDER		مالا	20161
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Project Manager:		- 11	5.2 5.1	() (1) Pestii	03/TKN)
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MA DEB MCB Data Enhancement Affirm to		DATA QUALITY OBJECTIVES	CHAIN-	CHAIN-OF-CUSTODY RECORD	
☐ YES ☐ NO MCP Data Certification required.	d. State Standard Collivoration	Project Specific QC		NOTE: All samples submitted subject to Standard Terms and Conditions on reverse hereof.	werse hereof.
☐ YES ☐ NO MCP Drinking Water Sample included. (Require collection of contingent duplicate sample.	□CT XMCP GW-1/S-1	specific QC. Projectis bend L. A includes require project specific QC includes Sample Duplicates, Matrix Spikes, and/or Matrix Spike Duplicates, Laboratory QC is not project specific unless prearranged. Project specific QC	Mushed by S	Time Received by:	Receipt Temperature:
Signature: U.A.A.D.+[MA Drinking Water	samples are charged on a per sample basis. Each MS, MSD and Sample Duplicate requires an additional sample aliquot	Relinquished by: Date	Time Received by:	Container Count:
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lyzed to 8271	liquots remaining	iting, dreet	1 22 d-10	Trip blanks are also required, if VOA sample collected)	☐ YES ☐ NO MCP Drinking Water Sample included (Require collection of contingent duplicate sample.)	☐ YES ☐ NO MCP Data Certification required.	MA DEP MCP Data Enhancement Affirmation	REMARKS / SPECIAL INSTRUCTIONS	1:30W2411-P2-L17-02-C	1 waan-la-111-02-B	_	-	-	-	_	-10-TH-68-11F6W	-10-11-69-11AES		-	52 W 2A11-P2-UT-01-A	SAMPLE		INSTHUCTIONS: Use separate line for each container (except replicates)) TRANDER	ager:	WILLAP		11-00 J	DETHEN STY	3000	LYTILAL	
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	☐ Matrix Spike Duplicate	Sample Duplicate Sample Duplicate		samples are charged on a per sample basis. Each MS, MSD and Sample Duplicate requires an additional sample aliquot		Many regulatory pro-	Project Specific QC	807 BÁTA QUALITY OBJECTIVES	- - - - - - - - - - -											Х	SOOML/16 or Plastic 1U/32 or Plastic 120mL Sterile HCI HNO, H,SO, NaOH Methanol Sodium Bisulfate ICE YES NO	Preservation Fillered	☐ GWA Quote:		☐ Purchase Order No.:	BILLING	☐ Please FAX to: KAT	ž	☐ PRIORITY (5 Business Days) ☐ RUSH (RAN-	STANDARD (10 Business Days)	TURNAROUND	CHAIN-OF-CUSTOD AND WORK ORDER	
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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:

Matrix:

Units:

EPA 8270C Modified

QC Batch ID: EP-2138-M

Soil ug/Kg Instrument ID: Extracted:

Analyzed:

MS-6 HP 6890 09-08-05 11:00 09-14-05 15:29 Instrument ID: Extracted:

MS-6 HP 6890 09-08-05 11:00 09-14-05 16:08

Analyst:

IIT

Analyzed: Analyst:

CAS Number	Analyte		LC	S		LC	S Duplicate		QCLim	its
	7.5	Spiked	Measured	Recovery	Spiked	Measured	Recovery	RPD	Spike	RPD
91-20-3	Naphthalene	330	190	58 %	330	210	64 %	10 %	40 - 140 %	20%
91-57-6	2-Methylnaphthalene	330	210	64 %	330	250	76 %	17 %	40 - 140 %	20%
85-01-8	Phenanthrene	330	270	82 %	330	270	82 %	0 %	40 - 140 %	20%
83-32-9	Acenaphthene	330	270	82 %	330	290	88 %	7 %	40 - 140 %	20%
208-96-8	Acenaphthylene	330	250	76 %	330	270	82 %	8 %	40 - 140 %	20%
86-73-7	Fluorene	330	260	79 %	330	280	85 %	7 %	40 - 140 %	20%
120-12-7	Anthracene	330	270	82 %	330	280	85 %	4 %	40 - 140 %	20%
206-44-0	Fluoranthene	330	310	94 %	330	330	100 %	6 %	40 - 140 %	20%
129-00-0	Pyrene	330	310	94 %	330	320	97 %	3 %	40 - 140 %	20%
56-55-3	Benzo[a]anthracene	330	310	94 %	330	330	100 %	6 %	40 - 140 %	20%
218-01-9	Chrysene	330	310	94 %	330	330	100 %	6 %	40 - 140 %	20%
205-99-2	Benzo[b]fluoranthene	330	300	91 %	330	310	94 %	3 %	40 - 140 %	20%
207-08-9	Benzo[k]fluoranthene	330	310	94 %	330	320	97 %	3 %	40 - 140 %	20%
50-32-8	Benzo[a]pyrene	330	300	91 %	330	310	94 %	3 %	40 - 140 %	20%
193-39-5	Indeno[1,2,3-c,d]pyrene	330	270	82 %	330	280	85 %	4 %	40 - 140 %	20%
53-70-3	Dibenzo[a,h]anthracene	330	250	76 %	330	260	79 %	4 %	40 - 140 %	20%
191-24-2	Benzo[g,h,i]perylene	330	270	82 %	330	290	88 %	7 %	40 - 140 %	20%

QC Surrogate Compound	Spiked	Measured	Recovery	Spiked	Measured	Recovery	QC Limits
ortho -Terphenyl	2,700	2,100	78 %	2,700	2,200	81 %	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:

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

QC Batch ID: Matrix: EP-2138-M Soil Instrument ID:

MS-6 HP 6890

Extracted: Analyzed: 09-08-05 11:00 09-14-05 17:27

40 - 140 %

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- <i>7</i> 3-7	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- <i>7</i> 0-3	Dibenzo[a,h]anthracene	BRL	ug/Kg	10
191-24-2	Benzo[g,h,i]perylene	BRL	ug/Kg	10

ortho-Terphenyl
Method Reference:

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

2700

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

2100

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:

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.



Quality Control Report Laboratory Control Samples

LCS

LCSD

Extracted:

Category:

MA DEP EPH Method

QC Batch ID: EP-2138-M

Matrix: Units:

Soil

mg/Kg

Instrument ID: Extracted:

Analyzed (AL):

Analyzed (AR):

Analyst:

GC-7 HP 5890

09-08-05 11:00

09-09-05 14:18

09-09-05 14:58

Instrument ID:

GC-7 HP 5890 09-08-05 11:00 09-09-05 15:42

Analyzed (AL): Analyzed (AR): 09-09-05 16:26

Analyst: MM

CAS Number	Analyte		1.0	IS .		- L(S Duplicate		QCLim	its
		Spiked	Measured	Recovery	Spiked	Measured	Recovery	RPD	Spike	RPD
111-84-2	n-Nonane (C ₉)	3.3	1.9	59 %	3.3	2.0	61 %	3 %	30 - 140 %	25%
124-18-5	n-Decane (C ₁₀)	3.3	2.2	68 %	3.3	2.3	70 %	3 %	40 - 140 %	25%
112-40-3	n-Dodecane (C ₁₂)	3.3	2.4	72 %	3.3	2.4	74 %	3 %	40 - 140 %	25%
629-59-4	n-Tetradecane (C ₁₄)	3.3	2.5	76 %	3.3	2.6	<i>7</i> 9 %	3 %	40 - 140 %	25%
544-76-3	n-Hexadecane (C ₁₆)	3.3	2.7	83 %	3.3	2.9	87 %	4 %	40 - 140 %	25%
593-45-3	n-Octadecane (C ₁₈)	3.3	2.9	89 %	3.3	3.1	93 %	5 %	40 - 140 %	25%
n/a	n-C9 to n-C18 Group	20	15	74 %	20	15	77 %	4 %	40 - 140 %	25%
629-92-5	л-Nonadecane (С ₁₉)	3.3	2.8	84 %	3.3	2.9	88 %	5 %	40 - 140 %	25%
112-95-8	n -Eicosane (C ₂₀)	3.3	2.9	89 %	3.3	3.1	94 %	6 %	40 - 140 %	25%
629-97-0	n-Docosane (C ₂₂)	3.3	2.9	89 %	3.3	3.1	94 %	5 %	40 - 140 %	25%
646-31-1	n-Tetracosane (C ₂₄)	3.3	2.9	89 %	3.3	3.1	95 %	6 %	40 - 140 %	25%
630-01-3	n-Hexacosane (C ₂₆)	3.3	2.9	87 %	3.3	3.0	92 %	5 %	40 - 140 %	25%
630-02-4	n-Octacosane (C ₂₈)	3.3	2.8	85 %	3.3	3.0	90 %	5 %	40 - 140 %	25%
638-68-6	n-Triacontane (C ₃₀)	3.3	2.8	84 %	3.3	2.9	88 %	5 %	40 - 140 %	25%
630-06-8	n-Hexatriacontane (C ₃₆)	3.3	2.4	72 %	3.3	2.5	75 %	3 %	40 - 140 %	25%
n/a	n-C19 to n-C36 Group	26	22	85 %	26	24	90 %	5 %	40 - 140 %	25%
91-20-3	Naphthalene	3.3	2.2	67 %	3.3	2.3	69 %	3 %	40 - 140 %	25%
91-57-6	2-Methylnaphthalene	3.3	2.4	73 %	3.3	2.5	76 %	3 %	40 - 140 %	25%
208-96-8	Acenaphthylene	3.3	2.5	76 %	3.3	2.6	79 %	4 %	40 - 140 %	25%
83-32-9	Acenaphthene	3.3	2.3	71 %	3.3	2.4	74 %	4 %	40 - 140 %	25%
86-73-7	Fluorene	3.3	2.5	75 %	3.3	2.6	78 %	4 %	40 - 140 %	25%
85-01-8	Phenanthrene	3.3	2.7	81 %	3.3	2.8	84 %	3 %	40 - 140 %	25%
120-12-7	Anthracene	3.3	2.8	86 %	3.3	2.9	89 %	3 %	40 - 140 %	25%
206-44-0	Fluoranthene	3.3	2.7	82 %	3.3	2.8	85 %	4 %	40 - 140 %	25%
129-00-0	Pyrene	3.3	2.7	81 %	3.3	2.8	84 %	4 %	40 - 140 %	25%
56-55-3	Benzo[a]anthracene	3.3	2.7	82 %	3.3	2.8	86 %	5 %	40 - 140 %	25%
218-01-9	Chrysene	3.3	2.7	83 %	3.3	2.9	87 %	5 %	40 - 140 %	25%
205-99-2	Benzo[b]fluoranthene	3.3	2.7	82 %	3.3	2.8	86 %	5 %	40 - 140 %	25%
207-08-9	Benzo[k]fluoranthene	3.3	2.7	81 %	3.3	2.8	85 %	5 %	40 - 140 %	25%
50-32-8	Benzo[a]pyrene	3.3	2.7	81 %	3.3	2.8	84 %	5 %	40 - 140 %	25%
193-39-5	Indeno[1,2,3-c,d]pyrene	3.3	2.9	87 %	3.3	3.0	91 %	5 %	40 - 140 %	25%
53-70-3	Dibenzo[a,h]anthracene	3.3	2.8	83 %	3.3	2.9	89 %	6 %	40 - 140 %	25%
191-24-2	Benzo[g,h,i]perylene	3.3	2.7	81 %	3.3	2.9	87 %	7 %	40 - 140 %	25%
n/a	PAH Group	56	45	80 %	56	47	83 %	4 %	40 - 140 %	25%

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

	Frac	ionation B	reakthrough Evalua	tion		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-2138-M

ortho-Terphenyl

Matrix:

Soil

Instrument ID:

GC-7 HP 5890

Extracted:

09-08-05 11:00

Analyzed (AL): Analyzed (AR): 09-09-05 14:21 09-09-05 15:05

40 - 140 %

Analyst:

MM

EPH Ranges	Concentration Notes	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	2.2	83 %	40 - 140 %
	2-Bromonaphthalene	2.7	2.0	76 %	40 - 140 %
Extraction:	Chloro-octadecane	2.7	2.0	75 %	40 - 140 %

Method Reference:

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

Sample extraction performed by microwave accelerated solvent extraction technique.

2.7

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.

86 %

- 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. ٥



Quality Control Report Laboratory Control Samples

LCS

LCSD

Category:

Matrix:

Units:

MA DEP EPH Method

QC Batch ID: EP-2140-M

Soil mg/Kg Instrument ID: Extracted: Analyzed (AL):

Analyzed (AR):

GC-7 HP 5890 09-09-05 15:00

09-12-05 15:40 09-12-05 16:25 Instrument ID: Extracted: Analyzed (AL):

GC-7 HP 5890 09-09-05 15:00 09-12-05 17:10

Analyzed (AR):

09-12-05 17:10 09-12-05 17:54

Analyst: MM Analyst: M

CAS Number	Analyte		LC.	'S	100	L. LC	S Duplicate	Alexandra (c.)	QC Lim	its
		Spiked	Measured	Recovery	Spiked	Measured	Recovery	RPD	Spike	RPD
111-84-2	л -Nonane (С ₉)	3.3	1.9	58 %	3.3	1.8	55 %	5 %	30 - 140 %	25%
124-18-5	n-Decane (C ₁₀)	3.3	2.3	69 %	3.3	2.1	65 %	6 %	40 - 140 %	25%
112-40-3	n-Dodecane (C ₁₂)	3.3	2.4	73 %	3.3	2.3	68 %	7 %	40 - 140 %	25%
629-59-4	n -Tetradecane (C ₁₄)	3.3	2.5	77 %	3.3	2.4	72 %	8 %	40 - 140 %	25%
544-76-3	n -Hexadecane (C ₁₆)	3.3	2.7	82 %	3.3	2.6	78 %	5 %	40 - 140 %	25%
593-45-3	n-Octadecane (C ₁₈)	3.3	2.9	88 %	3.3	2.8	84 %	4 %	40 - 140 %	25%
n/a	n-C9 to n-C18 Group	20	15	75 %	20	14	70 %	6 %	40 - 140 %	25%
629-92-5	n-Nonadecane (C ₁₉)	3.3	2.8	84 %	3.3	2.7	81 %	3 %	40 - 140 %	25%
112-95-8	n -Eicosane (C ₂₀)	3.3	2.8	86 %	3.3	2.7	83 %	4 %	40 - 140 %	25%
629-97-0	n-Docosane (C ₂₂)	3.3	2.8	84 %	3.3	2.7	81 %	4 %	40 - 140 %	25%
646-31-1	n -Tetracosane (C_{24})	3.3	2.8	85 %	3.3	2.7	82 %	3 %	40 - 140 %	25%
630-01-3	n -Hexacosane (С ₂₆)	3.3	2.7	82 %	3.3	2.6	79 %	3 %	40 - 140 %	25%
630-02-4	n -Octacosane (C ₂₈)	3.3	2.6	79 %	3.3	2.5	77 %	3 %	40 - 140 %	25%
638-68-6	n -Triacontane (C ₃₀)	3.3	2.6	78 %	3.3	2.5	76 %	3 %	40 - 140 %	25%
630-06-8	n -Hexatriacontane (C ₃₆)	3.3	2.3	71 %	3.3	2.3	69 %	3 %	40 - 140 %	25%
n/a	n-C19 to n-C36 Group	26	21	81 %	26	21	79 %	3 %	40 - 140 %	25%
91-20-3	Naphthalene	3.3	2.1	63 %	3.3	2.1	64 %	2 %	40 - 140 %	25%
91-57-6	2-Methylnaphthalene	3.3	2.3	69 %	3.3	2.4	72 %	4 %	40 - 140 %	25%
208-96-8	Acenaphthylene	3.3	2.4	73 %	3.3	2.5	75 %	2 %	40 - 140 %	25%
83-32-9	Acenaphthene	3.3	2.2	68 %	3.3	2.3	70 %	3 %	40 - 140 %	25%
86-73-7	Fluorene	3.3	2.4	72 %	3.3	2.5	76 %	4 %	40 - 140 %	25%
85-01-8	Phenanthrene	3.3	2.5	77 %	3.3	2.7	82 %	6 %	40 - 140 %	25%
120-12-7	Anthracene	3.3	2.7	83 %	3.3	2.9	87 %	5 %	40 - 140 %	25%
206-44-0	Fluoranthene	3.3	2.6	79 %	3.3	2.8	84 %	6 %	40 - 140 %	25%
129-00-0	Pyrene	3.3	2.6	77 %	3.3	2.7	82 %	6 %	40 - 140 %	25%
56-55-3	Benzo[a]anthracene	3.3	2.6	79 %	3.3	2.7	81 %	3 %	40 - 140 %	25%
218-01-9	Chrysene	3.3	2.7	82 %	3.3	2.7	82 %	0 %	40 - 140 %	25%
205-99-2	Benzo[b]fluoranthene	3.3	2.5	75 %	3.3	2.6	78 %	5 %	40 - 140 %	25%
207-08-9	Benzo[k]fluoranthene	3.3	2.6	78 %	3.3	2.6	78 %	0 %	40 - 140 %	25%
50-32-8	Benzo[a]pyrene	3.3	2.5	76 %	3.3	2.6	78 %	2 %	40 - 140 %	25%
193-39-5	Indeno[1,2,3-c,d]pyrene	3.3	2.6	79 %	3.3	2.7	81 %	3 %	40 - 140 %	25%
53-70-3	Dibenzo[a,h]anthracene	3.3	2.5	75 %	3.3	2.5	75 %	0 %	40 - 140 %	25%
191-24-2	Benzo[g,h,i]perylene	3.3	2.4	73 %	3.3	2.5	75 %	3 %	40 - 140 %	25%
n/a	PAH Group	56	42	75 %	56	44	78 %	3 %	40 - 140 %	25%

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

	Fract	ionation B	reakthrough 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-2140-M

Matrix:

Soil

Instrument ID:

GC-7 HP 5890

Extracted:

09-09-05 15:00

Analyzed (AL): Analyzed (AR): 09-12-05 14:11 09-12-05 14:56

Analyst:

MM

EPH Ranges	Concentration Notes	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 † °	BRL	mg/Kg	30
Unadjusted n-C11 to n-C22 Aromatic Hydrocarbons †	BRL	mg/Kg	30

QC Surrogate Co	ompound -	Spiked	Measured	Recovery	QC Limits
Fractionation:	2-Fluorobiphenyl	2.7	2.3	87 %	40 - 140 %
	2-Bromonaphthalene	2.7	2.3	88 %	40 - 140 %
Extraction:	Chloro-octadecane	2.7	2.0	76 %	40 - 140 %
	ortho-Terphenyl	2.7	2.1	79 %	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.
- ♦ n-C11 to n-C22 Aromatic Hydrocarbons range data excludes the method target analyte concentrations.



Quality Control Report Laboratory Control Samples

LCSD

Category: QC Batch ID: EP-2140-M

EPA 8270C Modified

Instrument ID: Extracted:

MS-6 HP 6890 09-09-05 15:00

JJT

Instrument ID: Extracted:

MS-6 HP 6890 09-09-05 15:00

40 - 140 %

Matrix: Units:

Soil ug/Kg Analyzed: Analyst:

09-13-05 03:58

Analyzed: Analyst:

09-13-05 04:37 JJT

	3 3			/	,, .		,,	24	,,,	
CAS Number	Analyte		LC	S		, LC	S Duplicate		QC Lim	its
		Spiked	Measured	Recovery	Spiked	Measured	Recovery	RPD	Spike	RPD
91-20-3	Naphthalene	330	200	61 %	330	210	64 %	5 %	40 - 140 %	20%
91-57-6	2-Methylnaphthalene	330	220	67 %	330	240	73 %	9 %	40 - 140 %	20%
85-01-8	Phenanthrene	330	240	73 %	330	250	76 %	4 %	40 - 140 %	20%
83-32-9	Acenaphthene	330	270	82 %	330	280	85 %	4 %	40 - 140 %	20%
208-96-8	Acenaphthylene	330	250	76 %	330	260	79 %	4 %	40 - 140 %	20%
86-73-7	Fluorene	330	250	76 %	330	260	79 %	4 %	40 - 140 %	20%
120-12-7	Anthracene	330	240	73 %	330	250	76 %	4 %	40 - 140 %	20%
206-44-0	Fluoranthene	330	270	82 %	330	290	88 %	7 %	40 - 140 %	20%
129-00-0	Pyrene	330	270	82 %	330	290	88 %	7 %	40 - 140 %	20%
56-55-3	Benzo[a]anthracene	330	270	82 %	330	290	88 %	7 %	40 - 140 %	20%
218-01-9	Chrysene	330	270	82 %	330	290	88 %	7 %	40 - 140 %	20%
205-99-2	Benzo[b]fluoranthene	330	260	79 %	330	280	85 %	7 %	40 - 140 %	20%
207-08-9	Benzo[k]fluoranthene	330	270	82 %	330	290	88 %	7 %	40 - 140 %	20%
50-32-8	Benzo[a]pyrene	330	260	79 %	330	290	88 %	11 %	40 - 140 %	20%
193-39-5	Indeno[1,2,3-c,d]pyrene	330	260	79 %	330	280	85 %	7 %	40 - 140 %	20%
53-70-3	Dibenzo[a,h]anthracene	330	240	73 %	330	260	79 %	8 %	40 - 140 %	20%
191-24-2	Benzo[g,h,i]perylene	330	270	82 %	330	290	88 %	7 %	40 - 140 %	20%
QC Surrogate	Compound	Spiked	Measured	Recovery	Spiked	Measured	Recovery	respect	QC Lim	ts

ortho-Terphenyl Method Reference:

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

70 %

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

2,700 2,100

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.

2,700 1,900

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

Instrument ID:

MS-6 HP 6890

Matrix:

EP-2140-M Soil Extracted: Analyzed:

09-09-05 15:00 09-13-05 03:19

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- <i>7</i>	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	2,300	86 %	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:

L 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