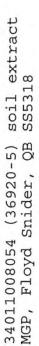
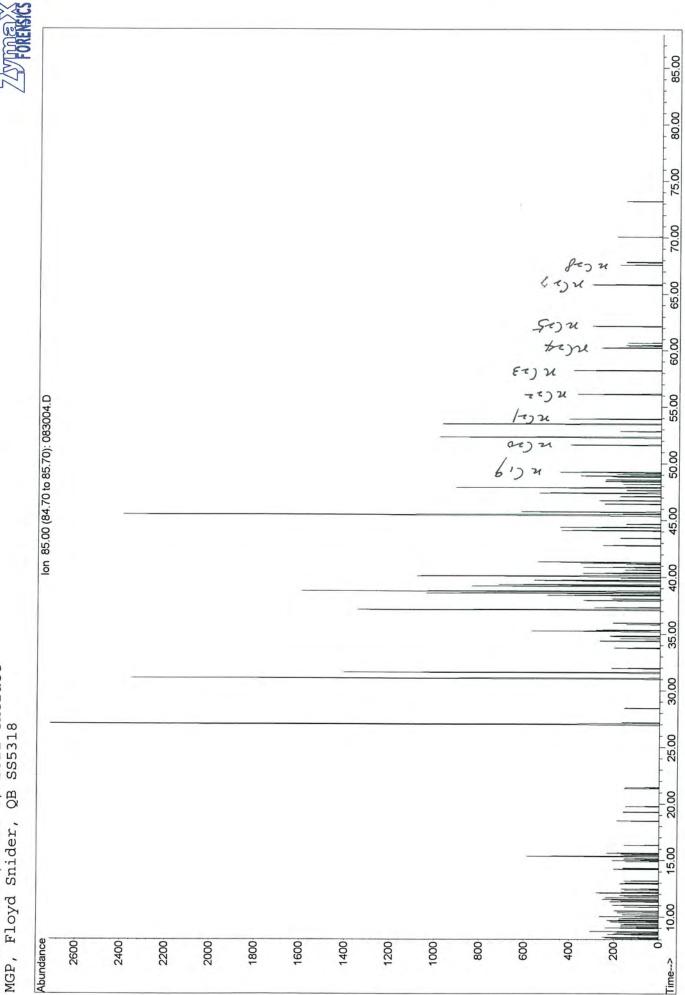
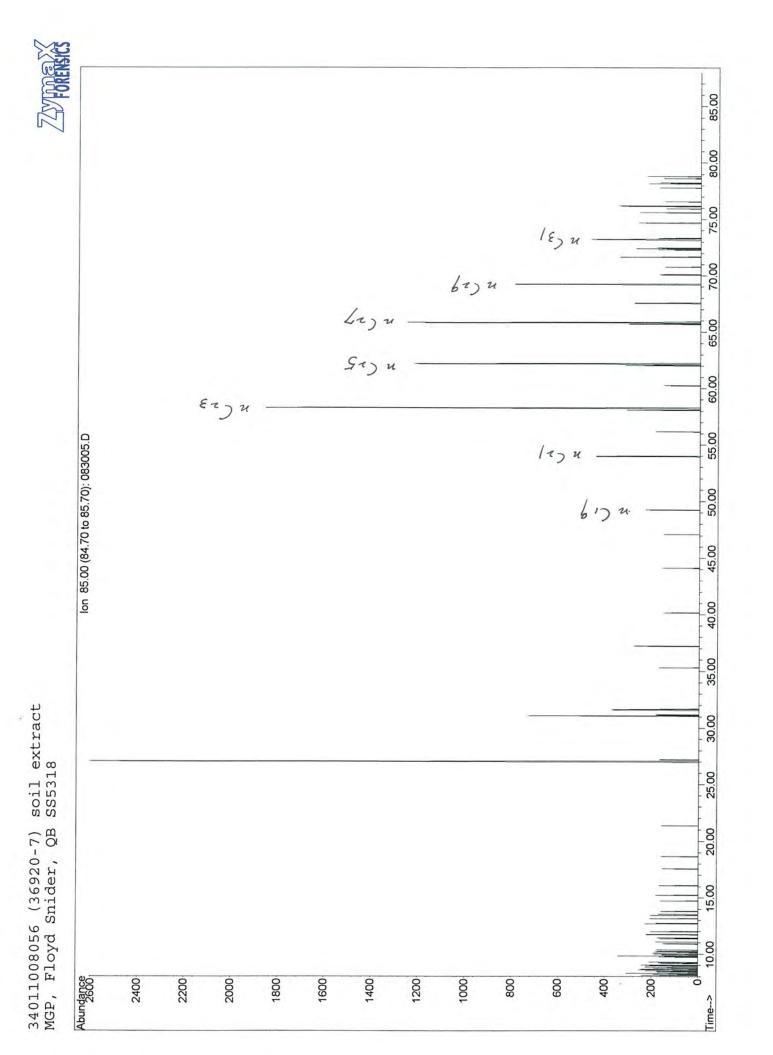
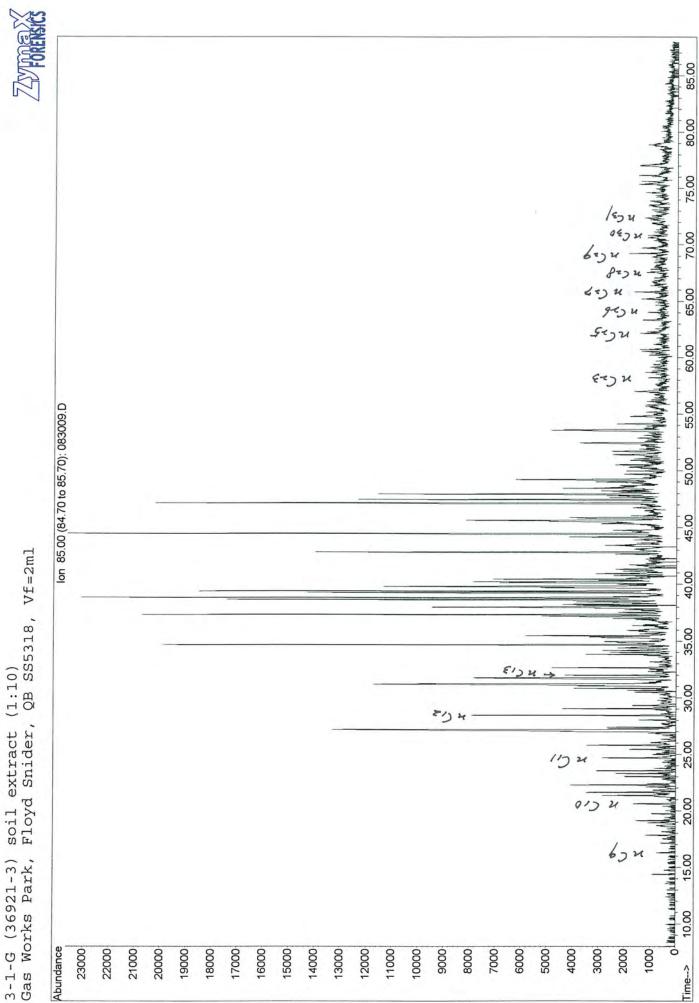


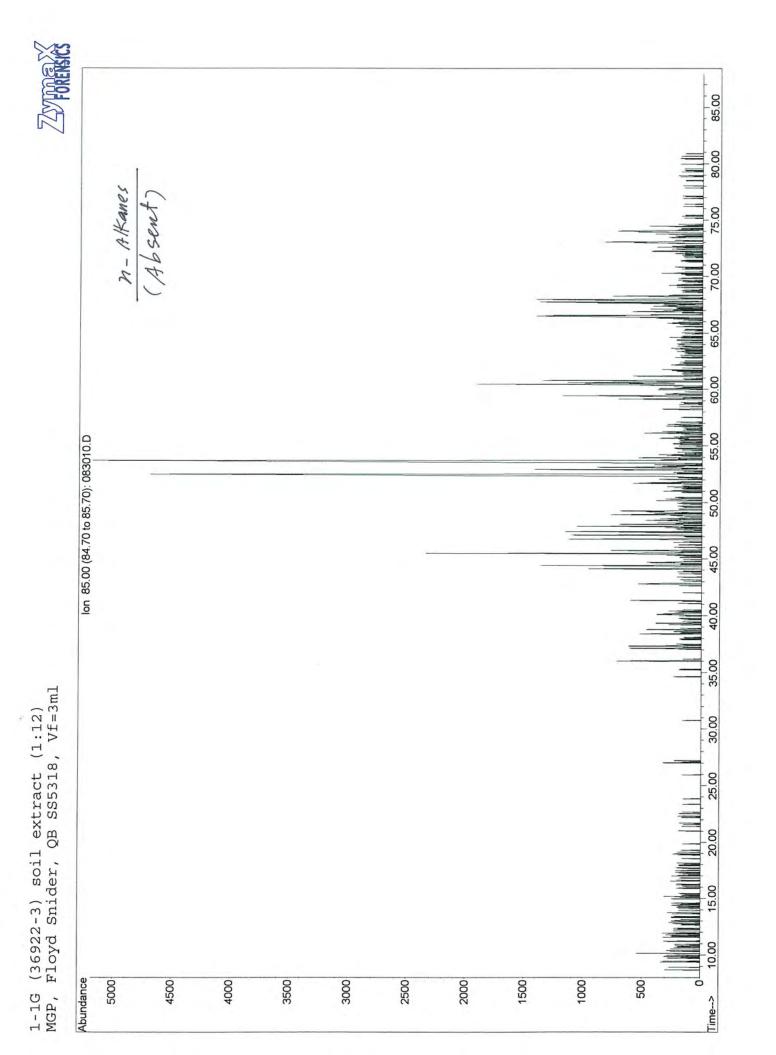
soil extract (1:7)
SS5318, Vf=3ml 34011008050 (36920-1) MGP, Floyd Snider, QB





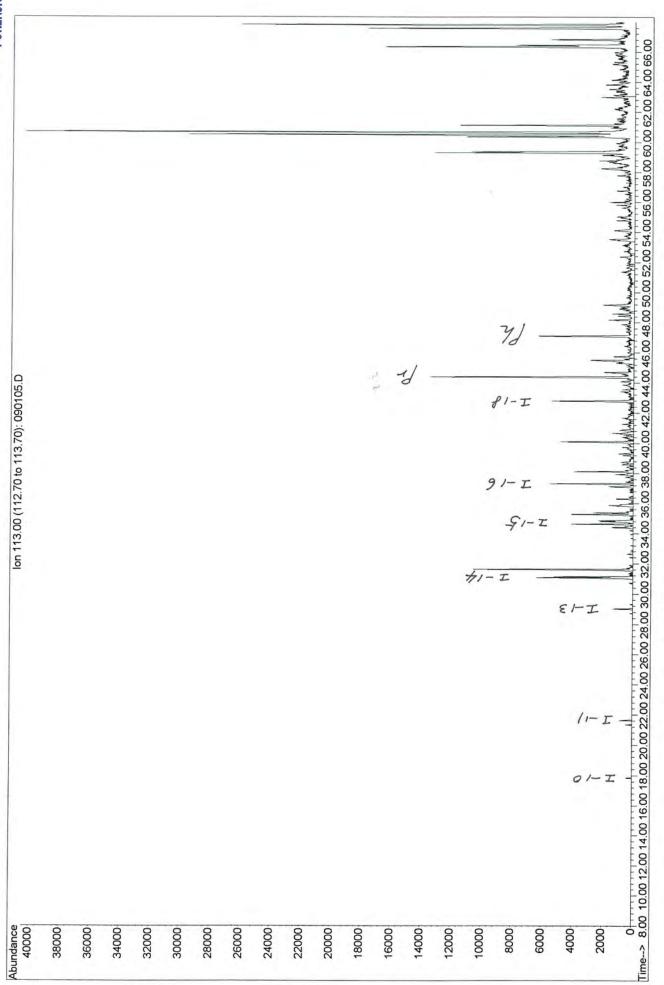






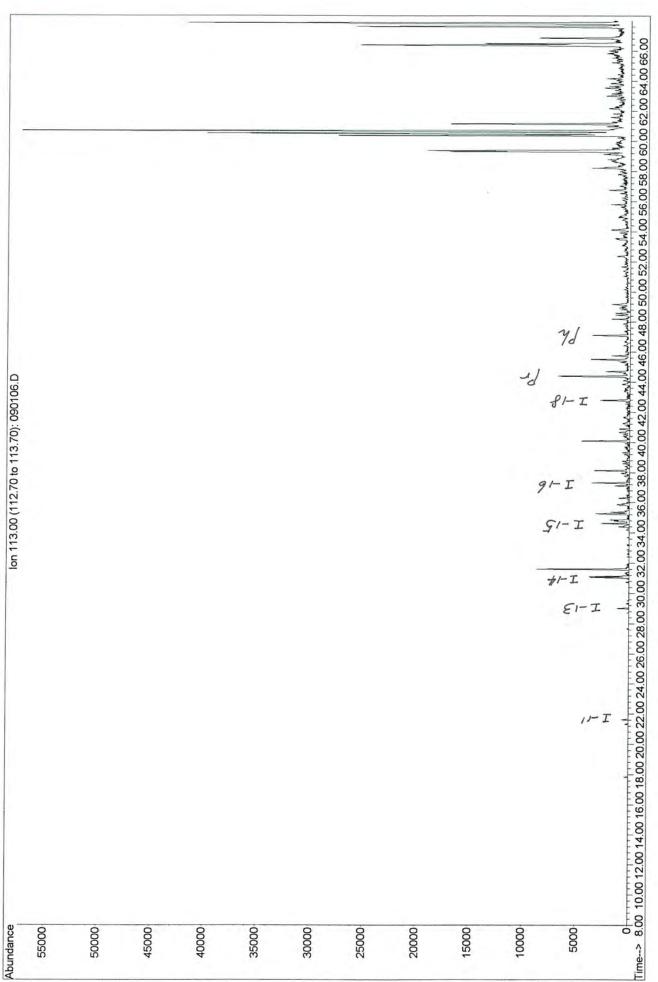
34011008001 (36917-1) soil extract (1:2) MGP, Floyd Snider, QB SS5305, Vf=1ml

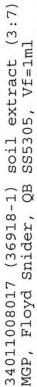


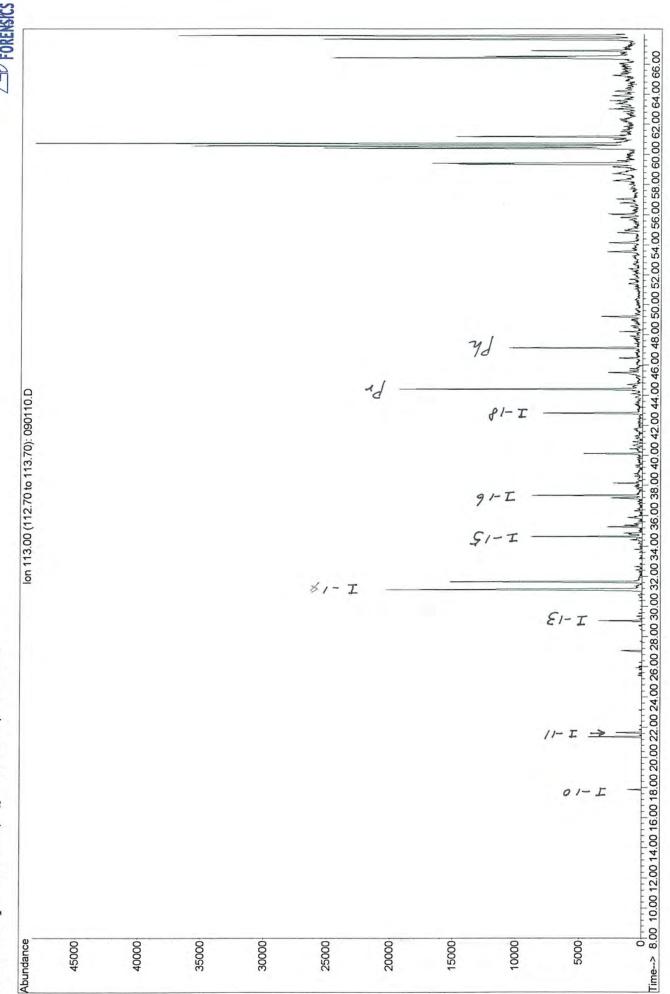


34011008011 (36917-11) soil extract (1:2) MGP, Floyd Snider, QB SS5305, Vf=1ml

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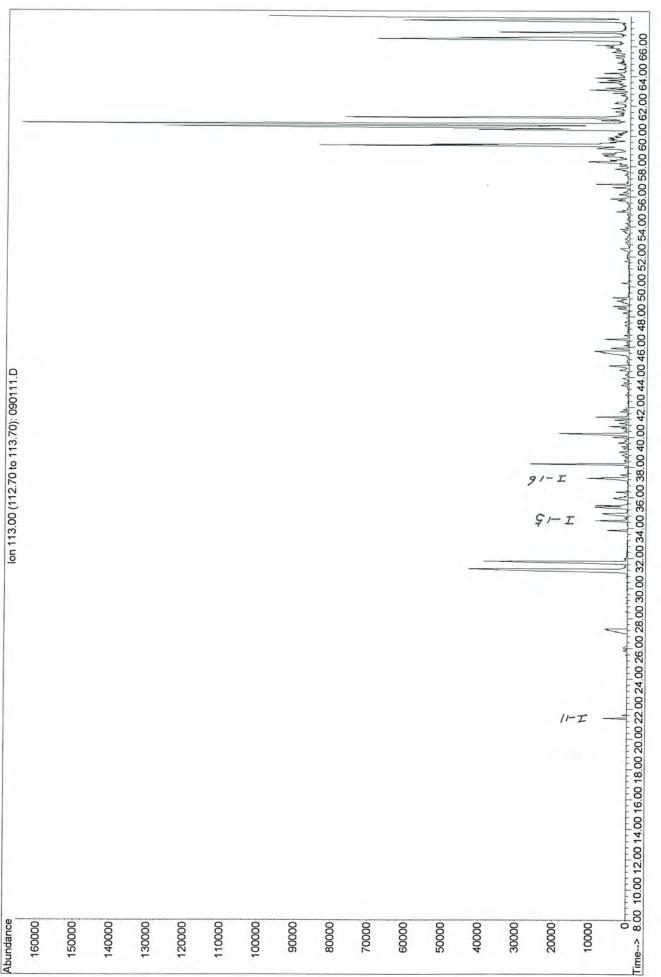




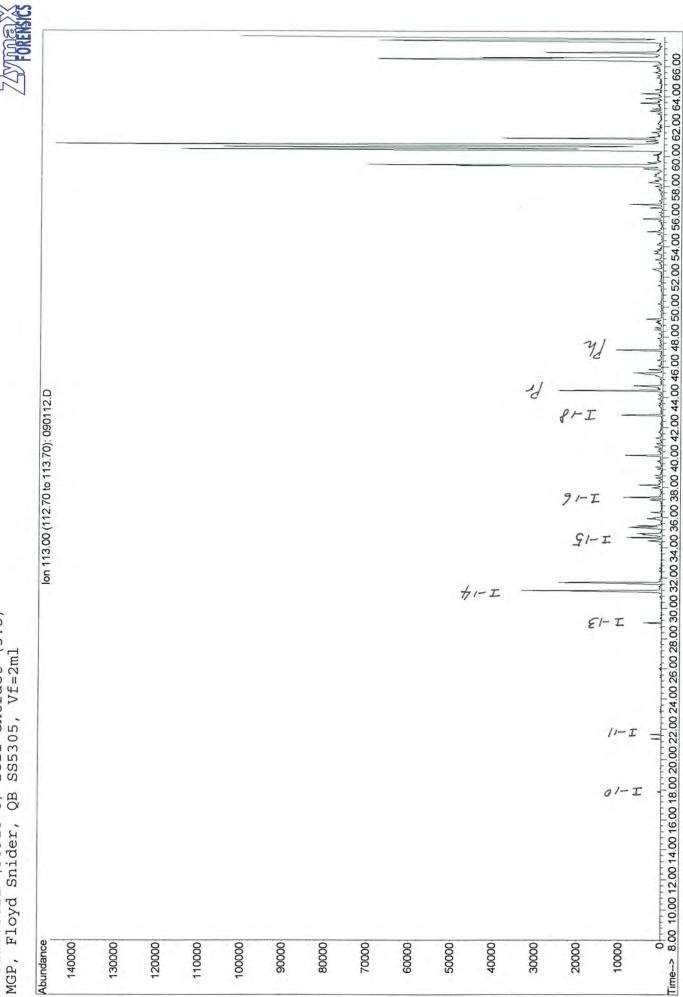


34011008020 (36918-4) soil extract (1:1.2) MGP, Floyd Snider, QB SS5305, Vf=2ml





soil extract (3:8) SS5305, Vf=2ml 34011008022 (36918-6) MGP, Floyd Snider, QB



34011008028 (36918-12) soil extract (1:1))

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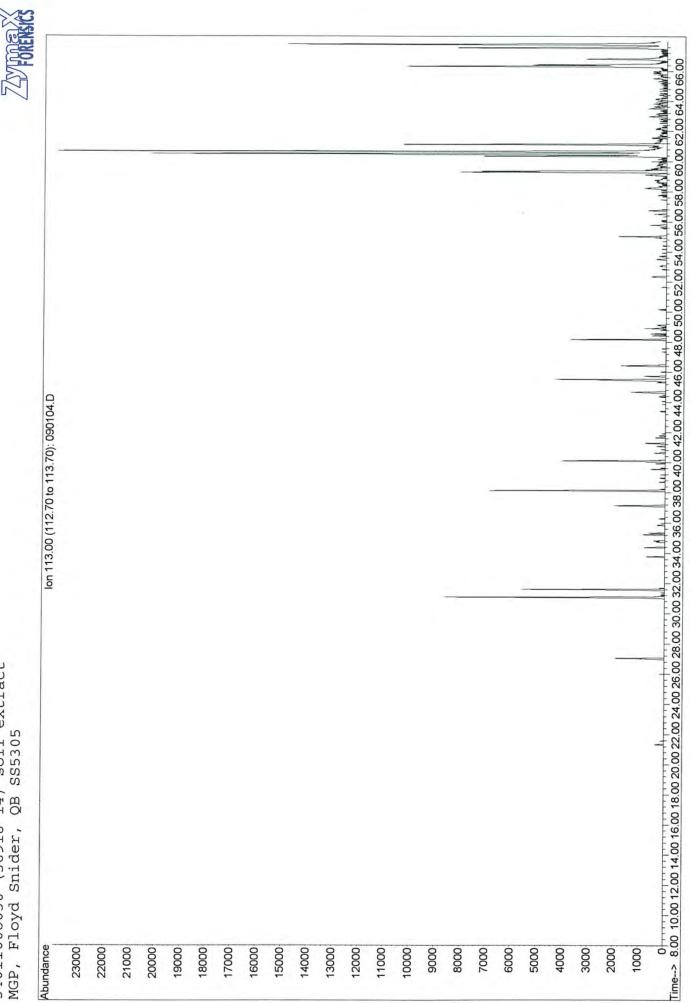
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34011008030 (36918-14) soil extract MGP, Floyd Snider, QB SS5305



34011008032 (36918-16) soil ext (3.5:6.5) MGP, Floyd Snider, QB SS5305, Vf=4ml

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34011008035 (36	Floyd Snide

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34011008040 (36919-4) soil extract (1:2) MGP, Floyd Snider, QB SS5318

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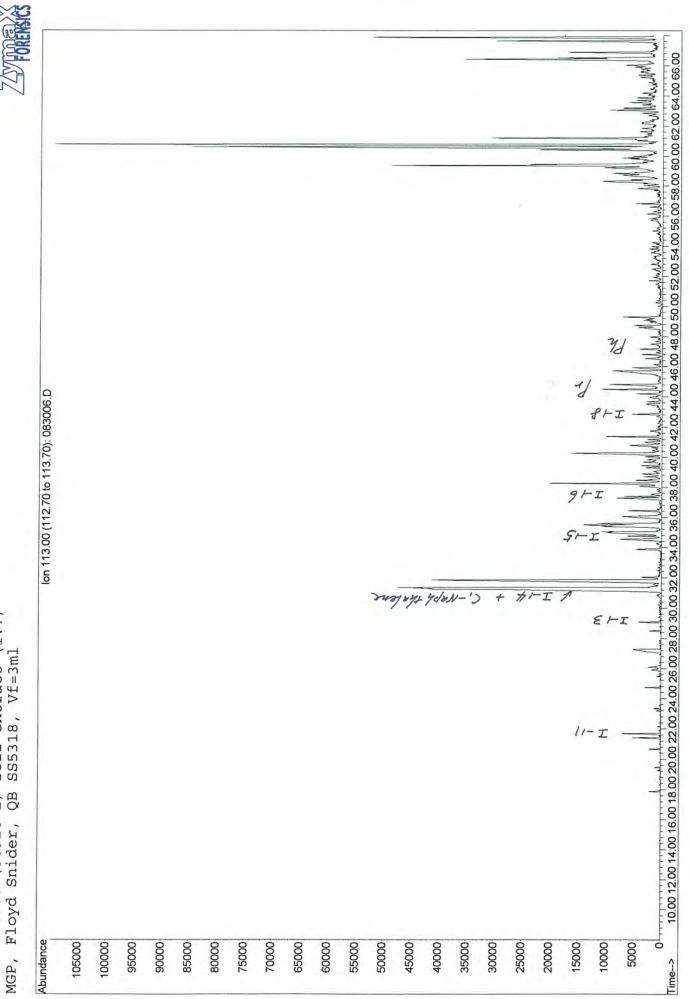
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34011008043 (36919-7) soil extract (1:2) MGP, Floyd Snider, QB SS5318, Vf=3ml

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soil extract (1:7)
SS5318, Vf=3ml 34011008050 (36920-1) MGP, Floyd Snider, QB



34011008054 (36920-5) soil extract MGP, Floyd Snider, QB SS5318

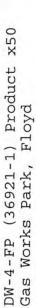
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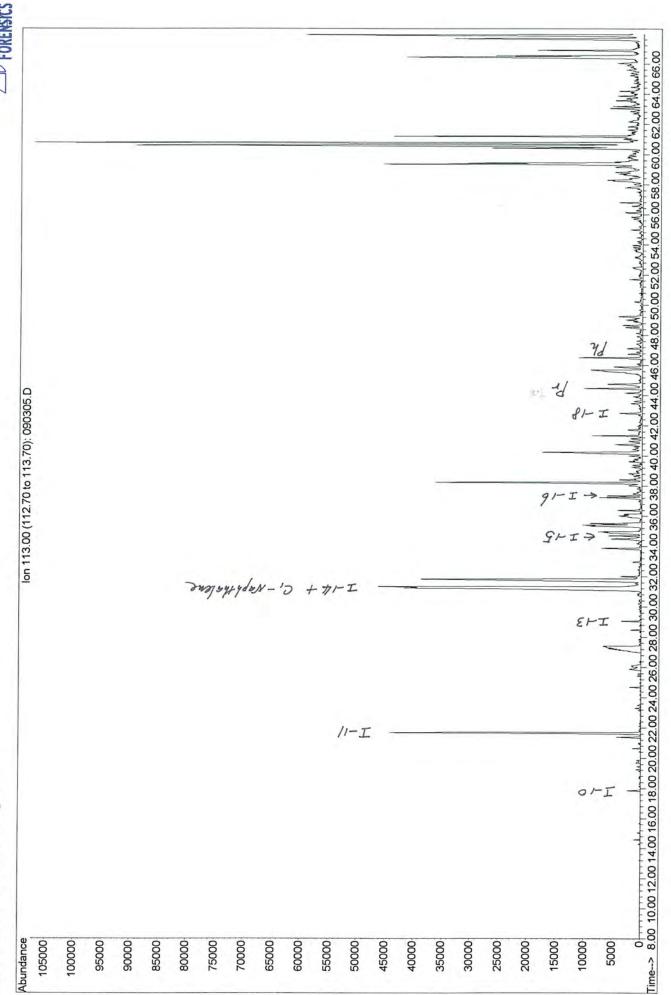
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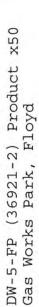
34011008056 (36920-7) soil extract MGP, Floyd Snider, QB SS5318

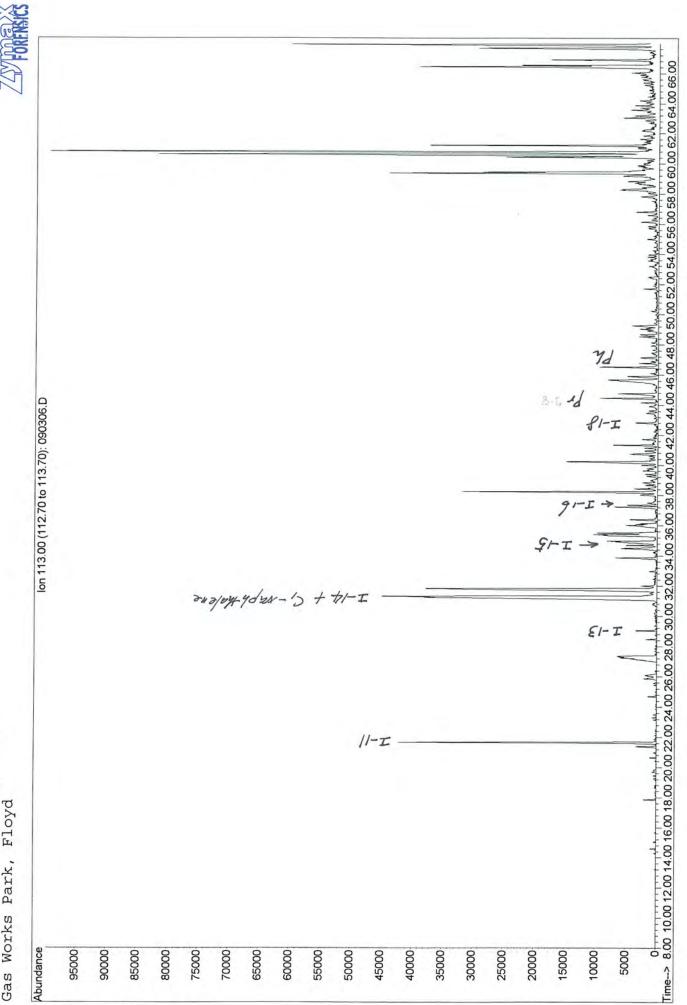


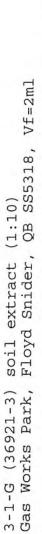
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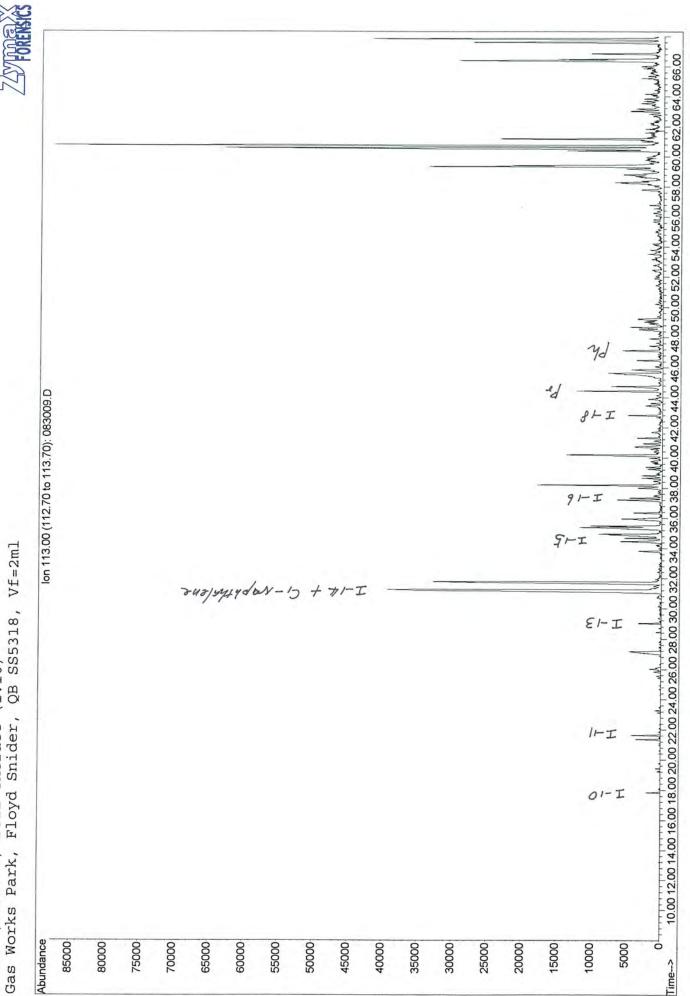












1-1G (36922-3) soil extract (1:12) MGP, Floyd Snider, QB SS5318, Vf=3ml

ZZVFRIRKS

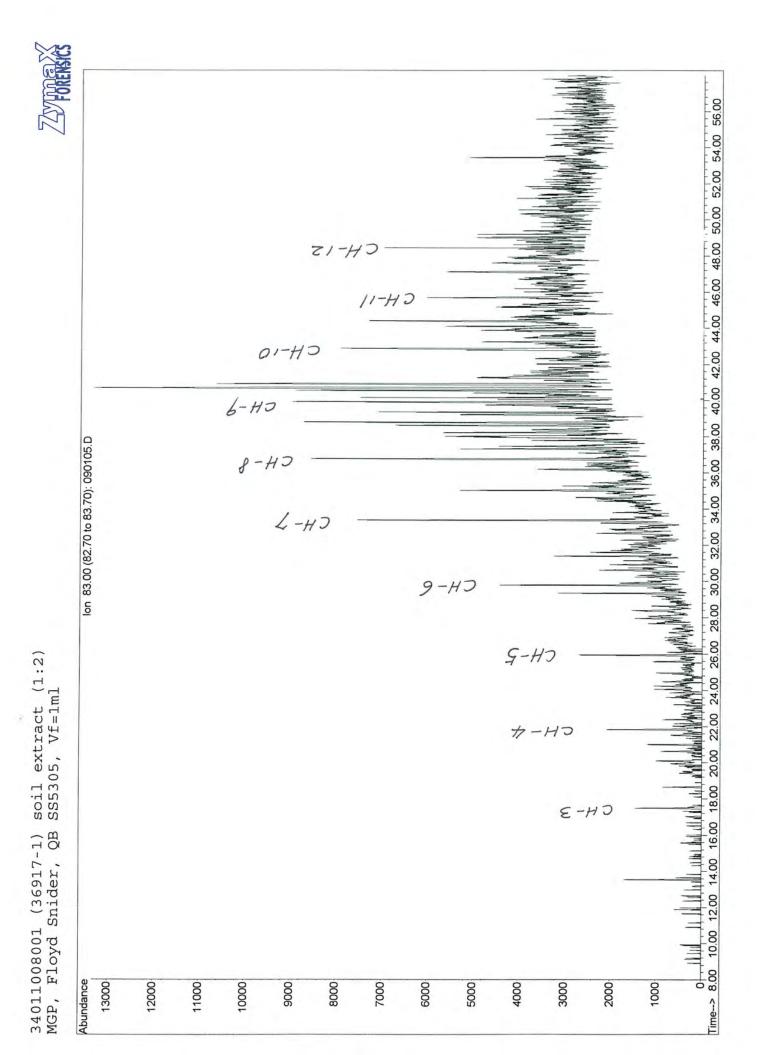
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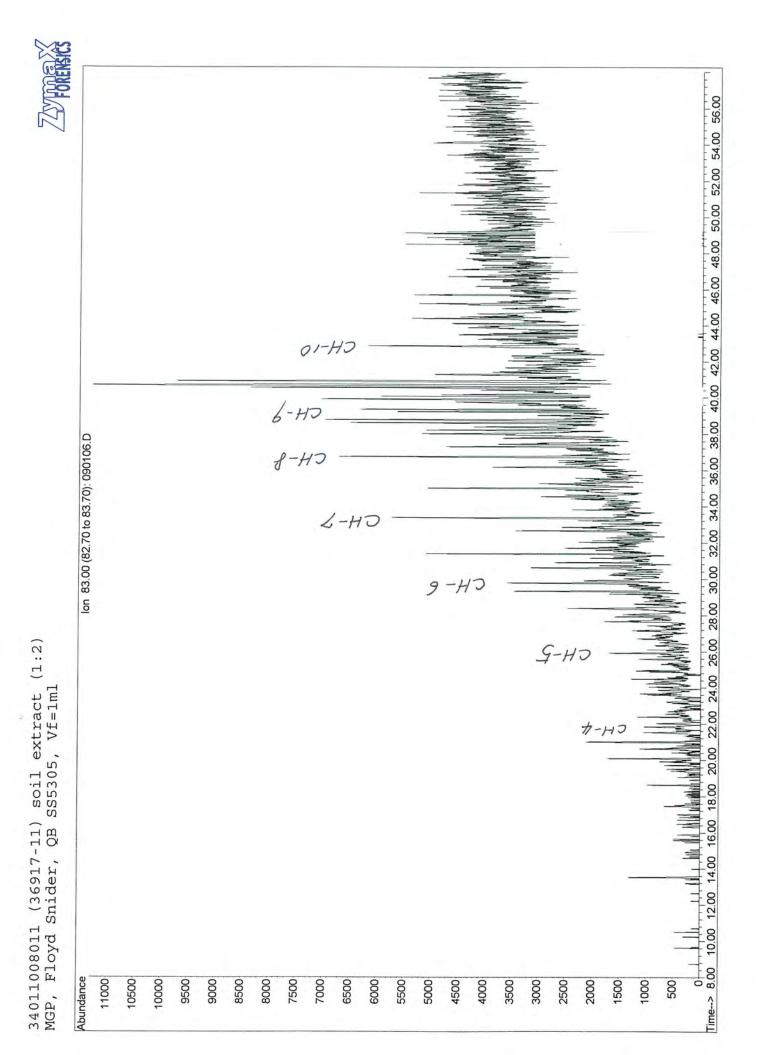


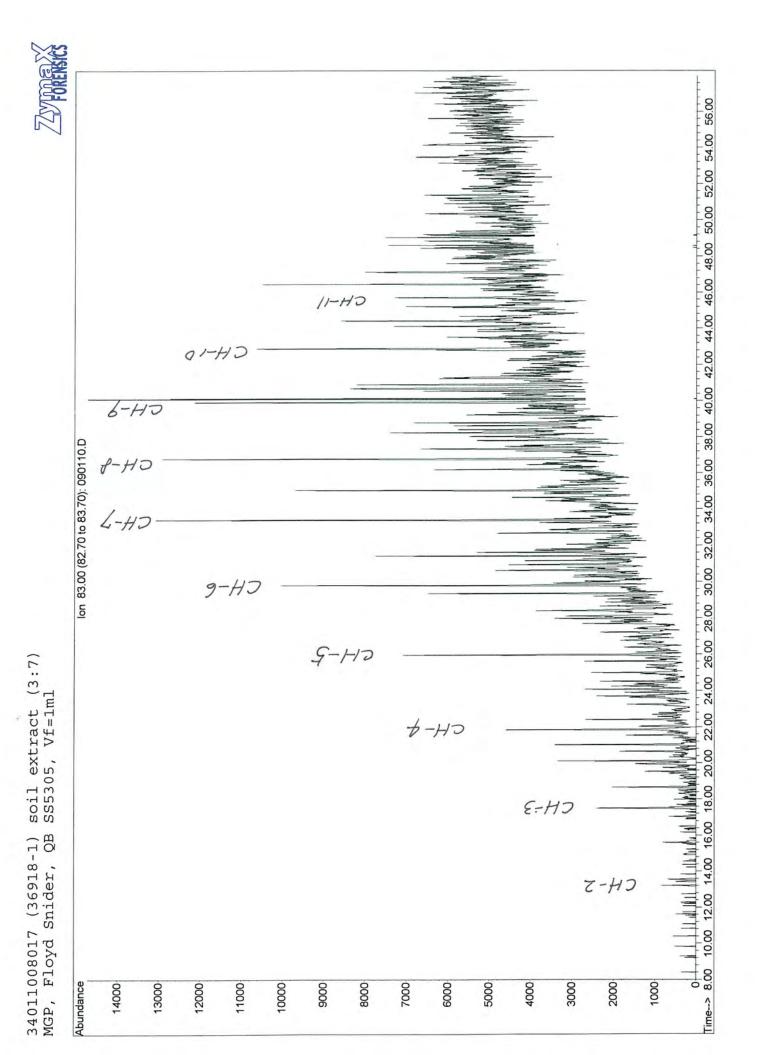
Table

Key for Alkylcyclohexanes at m/z 83

-	Symbol	Detail
	CH-1:	Methylcyclohexane
	CH-2:	Ethylcyclohexane
	CH-3:	Propylcylohexane
	CH-4:	Butylcyclohexane
	CH-5:	Pentylcyclohexane
	CH-6:	Hexylcyclohexane
	CH-7:	Heptylcyclohexane
	CH-8:	Octylcyclohexane
	CH-9:	Nonylcyclohexane
	CH-10:	Decylcyclohexane
	CH-11:	Undecylcyclohexane
à	CH-12:	Dodecylcyclohexane
	CH-13:	Tridecylcyclohexane
	CH-14:	Tetradecylcyclohexane



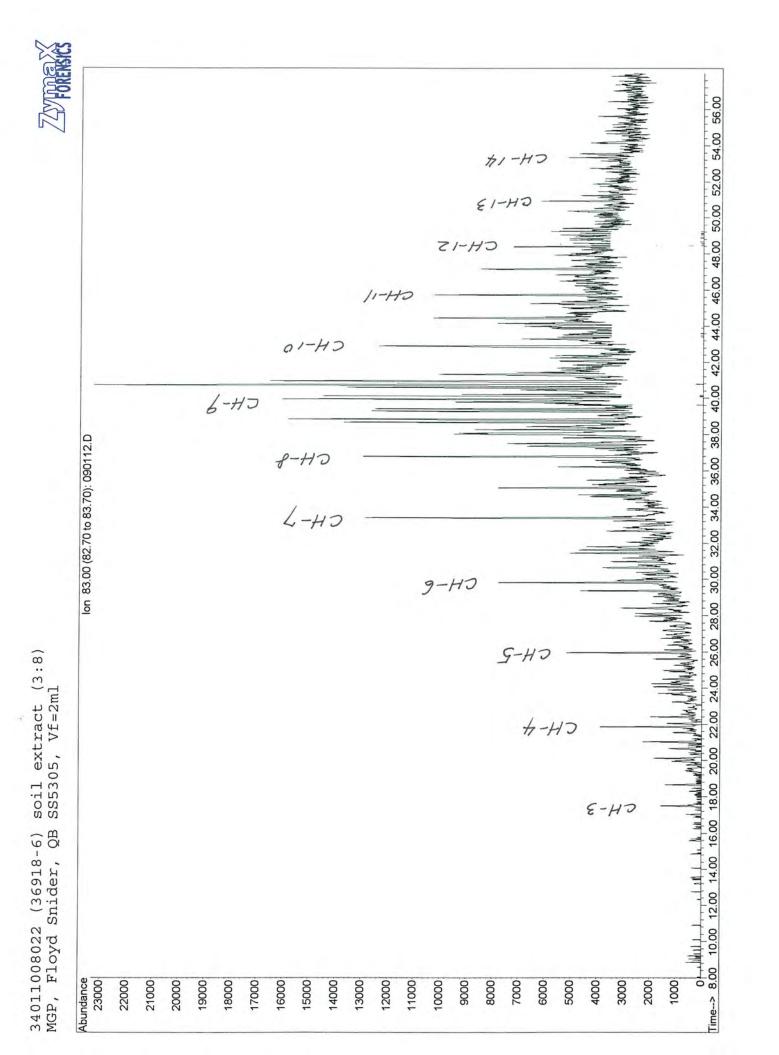


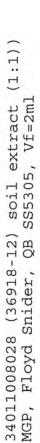


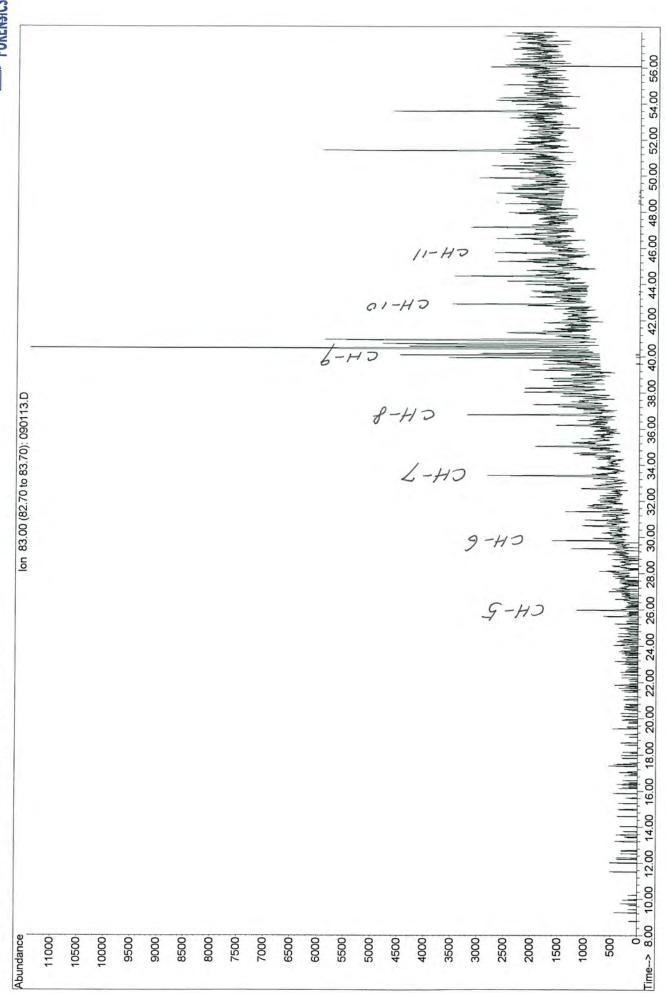
34011008020 (36918-4) soil extract (1:1.2) MGP, Floyd Snider, QB SS5305, Vf=2ml



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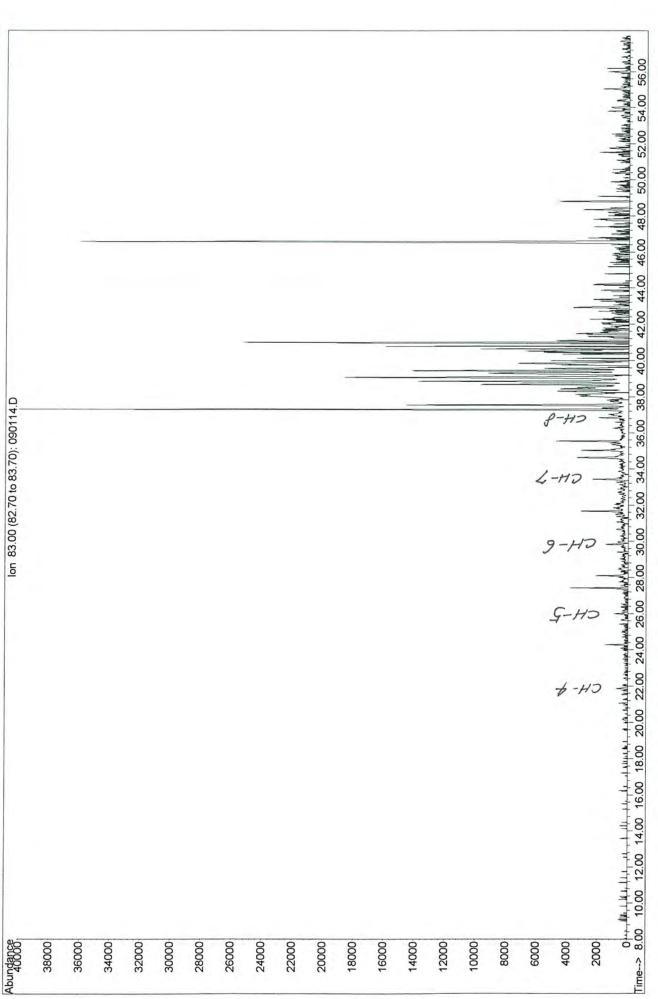
34011008030 (36918-14) soil extract MGP, Floyd Snider, QB SS5305

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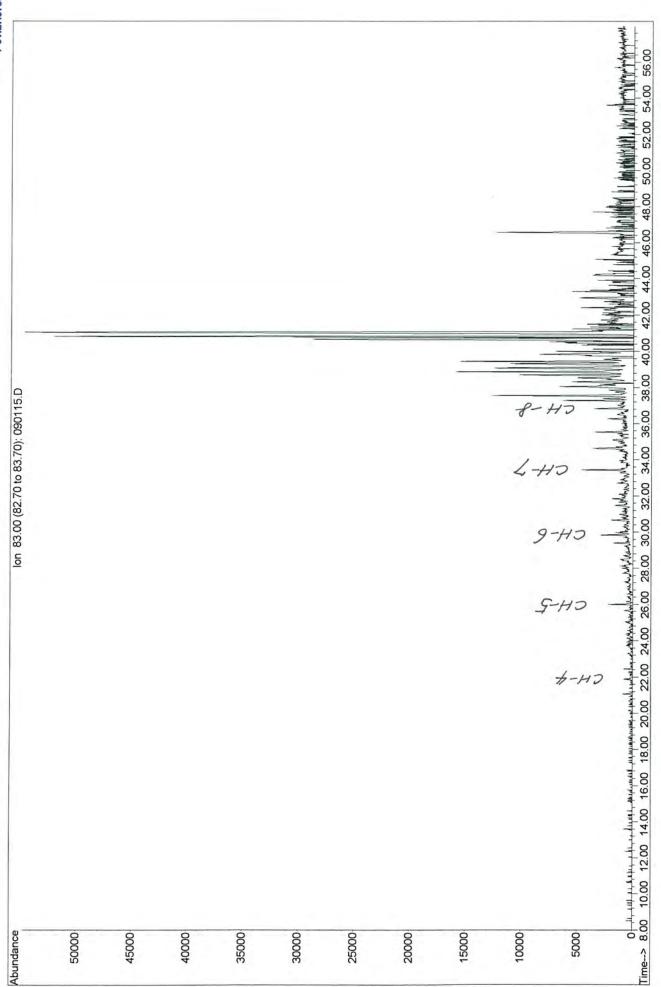
34011008032 (36918-16) soil ext (3.5:6.5) MGP, Floyd Snider, QB SS5305, Vf=4ml





34011008035 (36918-19) soil ext (4:6) MGP, Floyd Snider, QB SS5305, Vf=1ml

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34011008040 (36919-4) MGP, Floyd Snider, QB

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(1:2)	
soil extract (1:2) SS5318	

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34011008043 (36919-7) soil extract (1:2) MGP, Floyd Snider, QB SS5318, Vf=3ml

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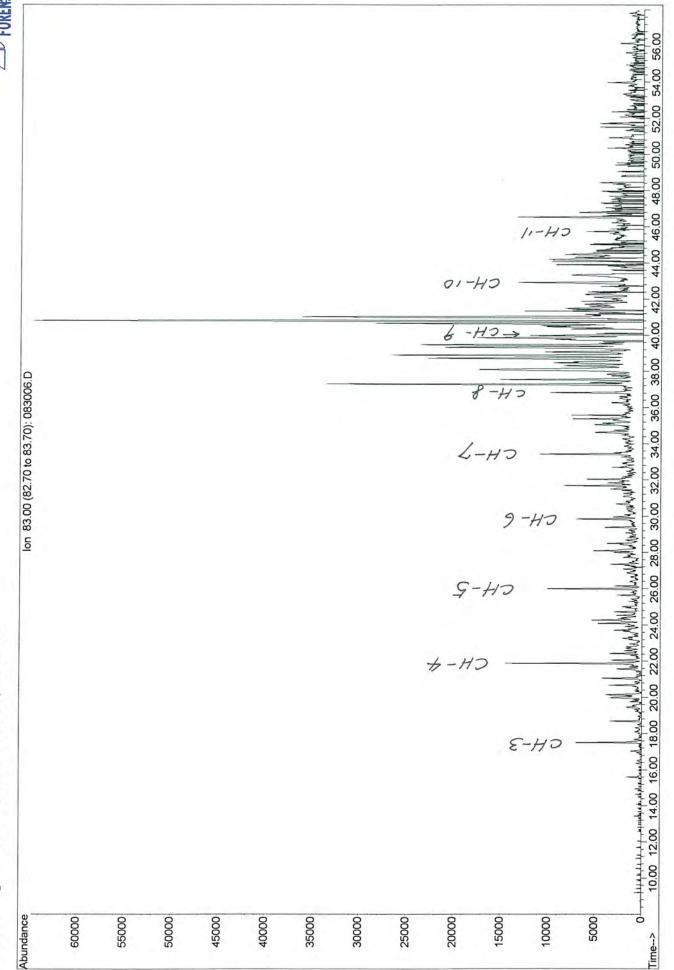
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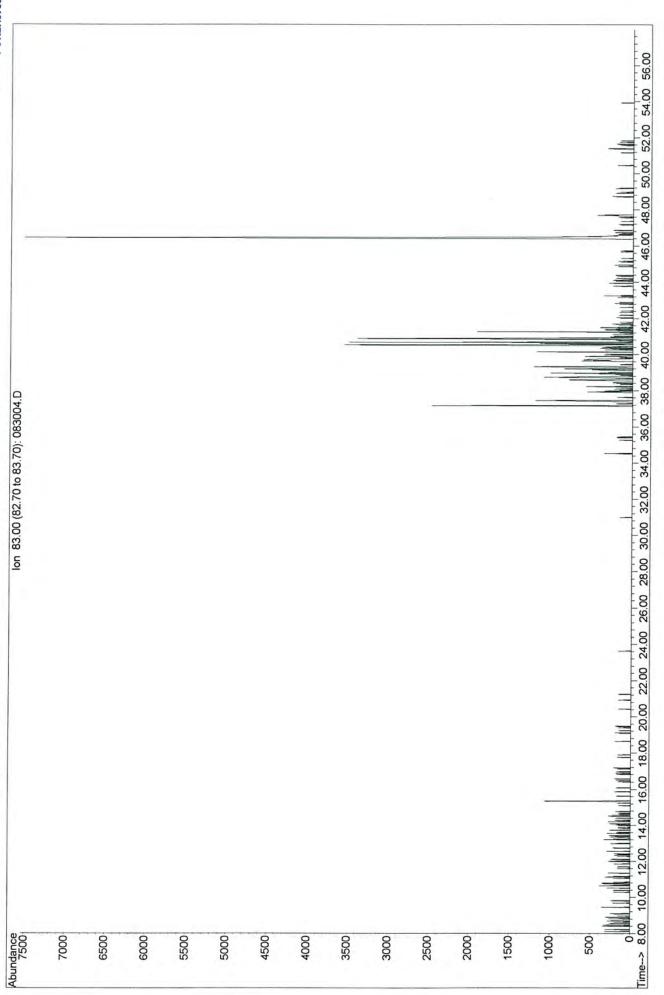
ZYPRIRICS

34011008050 (36920-1) soil extract (1:7) MGP, Floyd Snider, QB SS5318, Vf=3ml



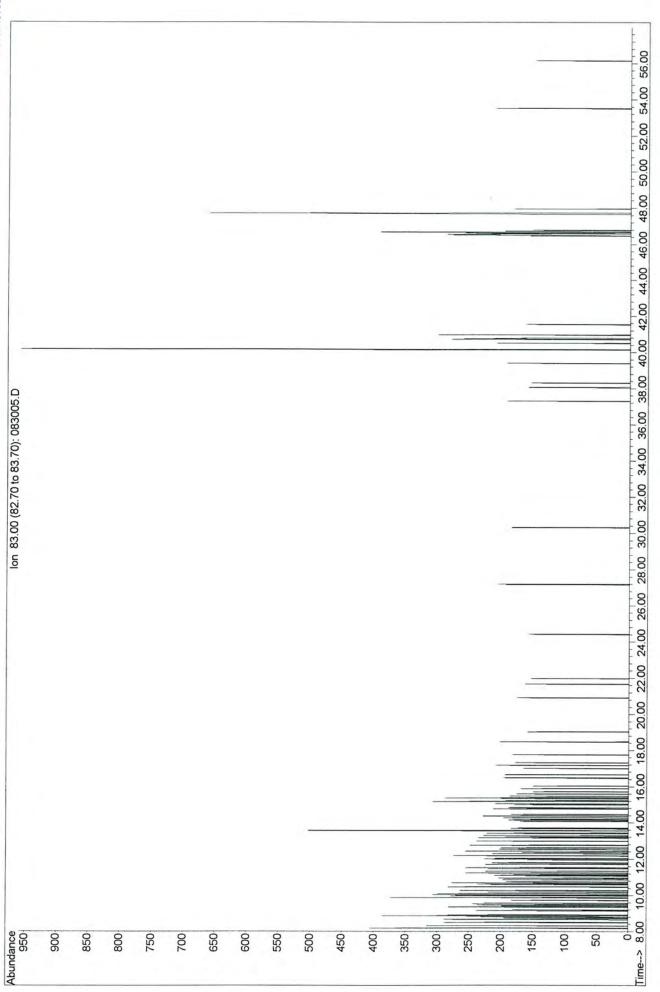
34011008054 (36920-5) soil extract MGP, Floyd Snider, QB SS5318

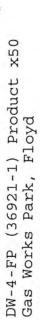


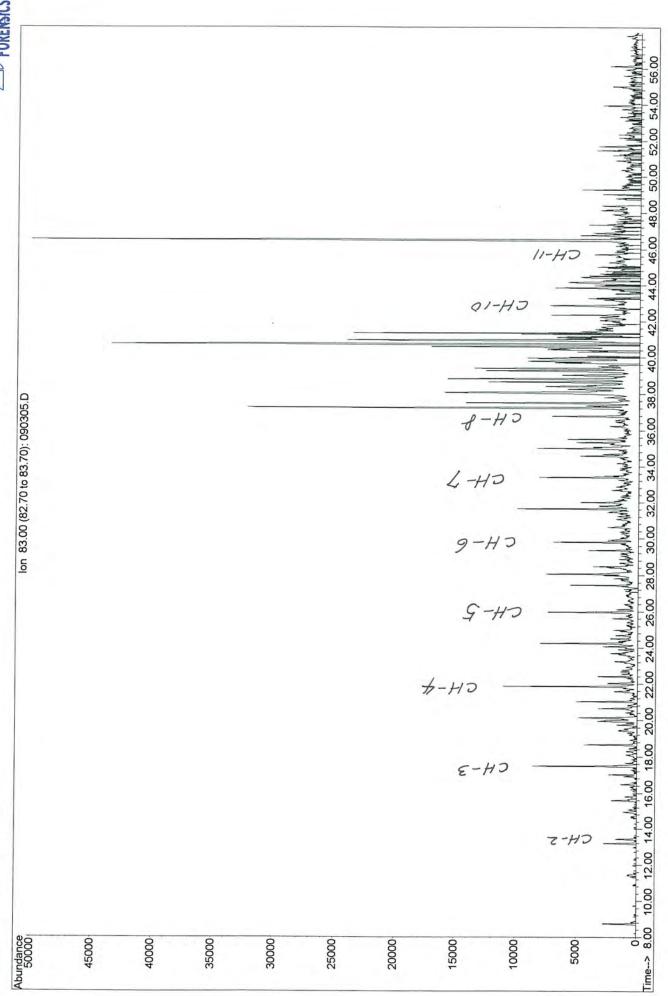


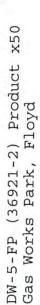
34011008056 (36920-7) soil extract MGP, Floyd Snider, QB SS5318

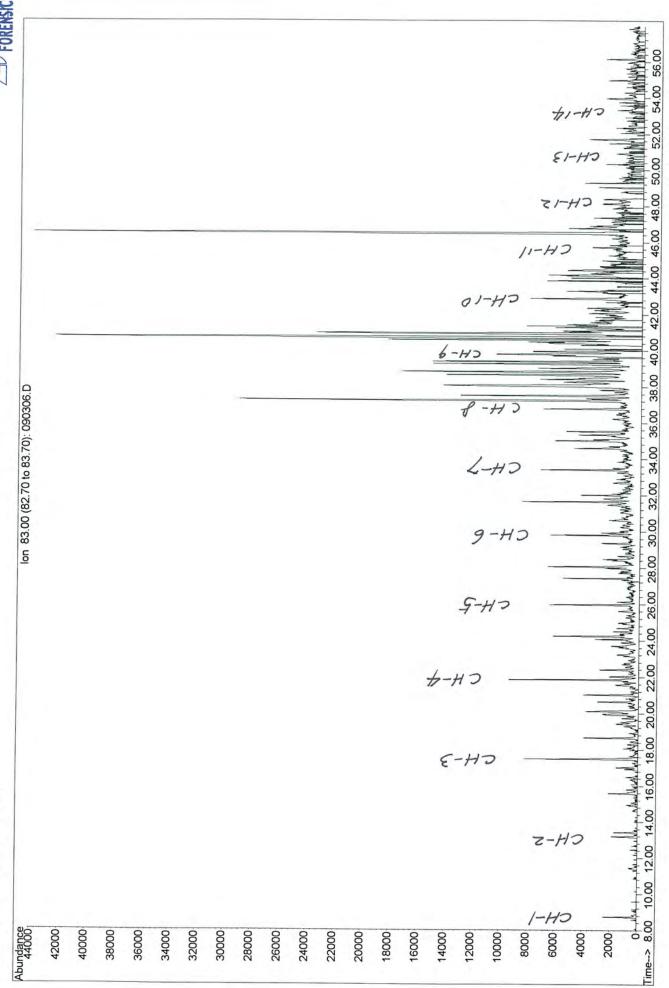




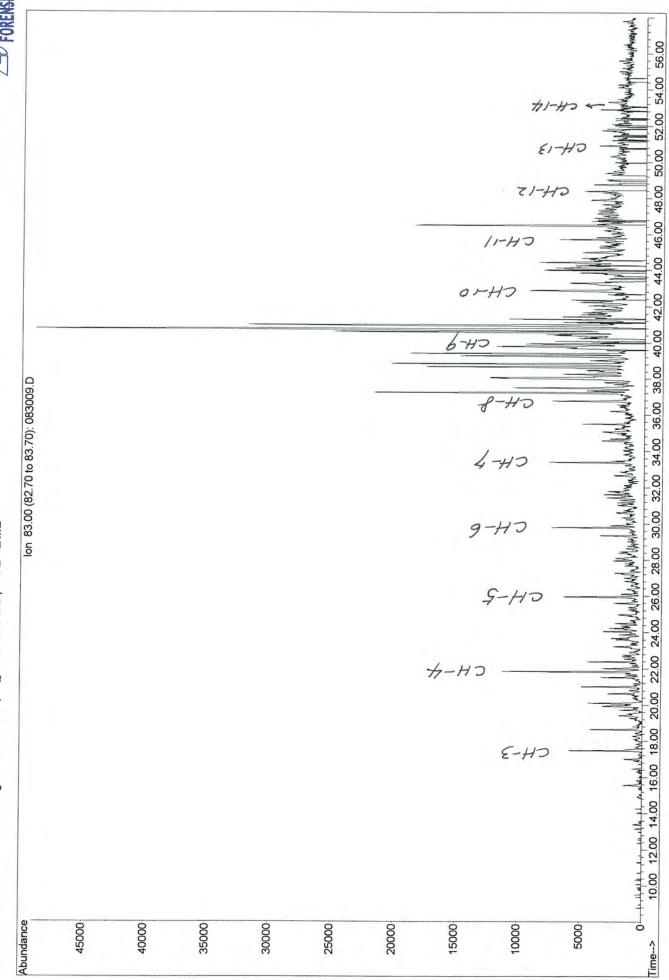






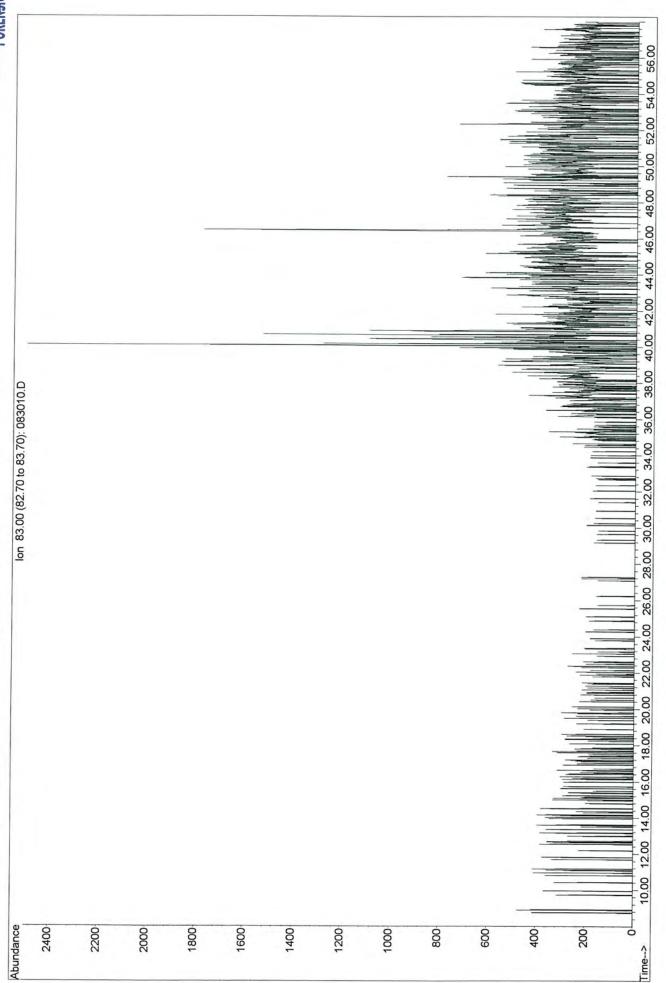


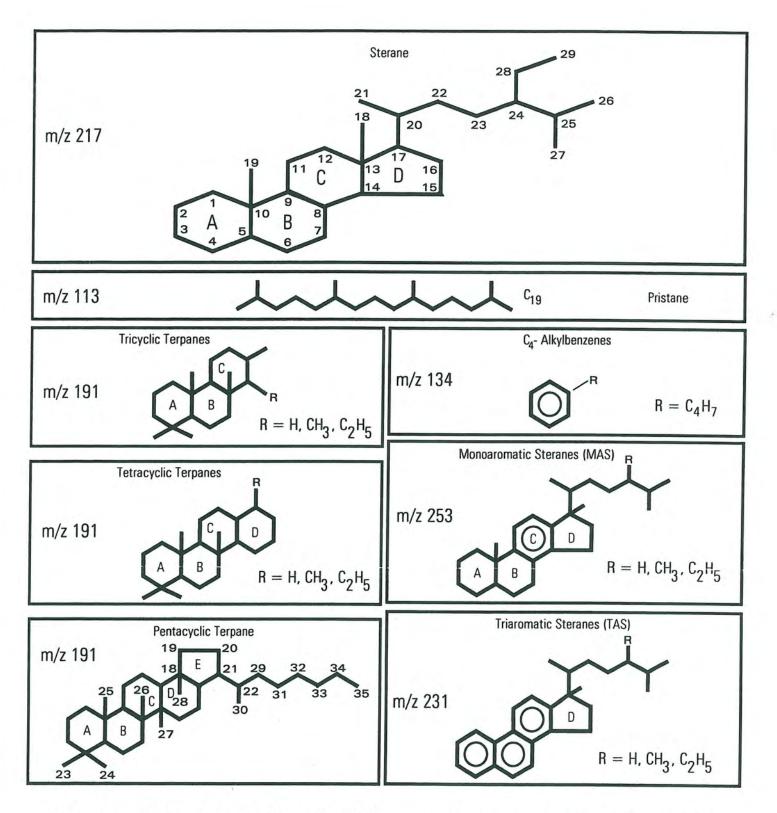
3-1-G (36921-3) soil extract (1:10) Gas Works Park, Floyd Snider, QB SS5318, Vf=2ml



1-1G (36922-3) soil extract (1:12) MGP, Floyd Snider, QB SS5318, Vf=3ml







The compound structures of pristane, C₄-alkylbenzenes, sterane; terpanes; monoaromatic and triaromatic steranes

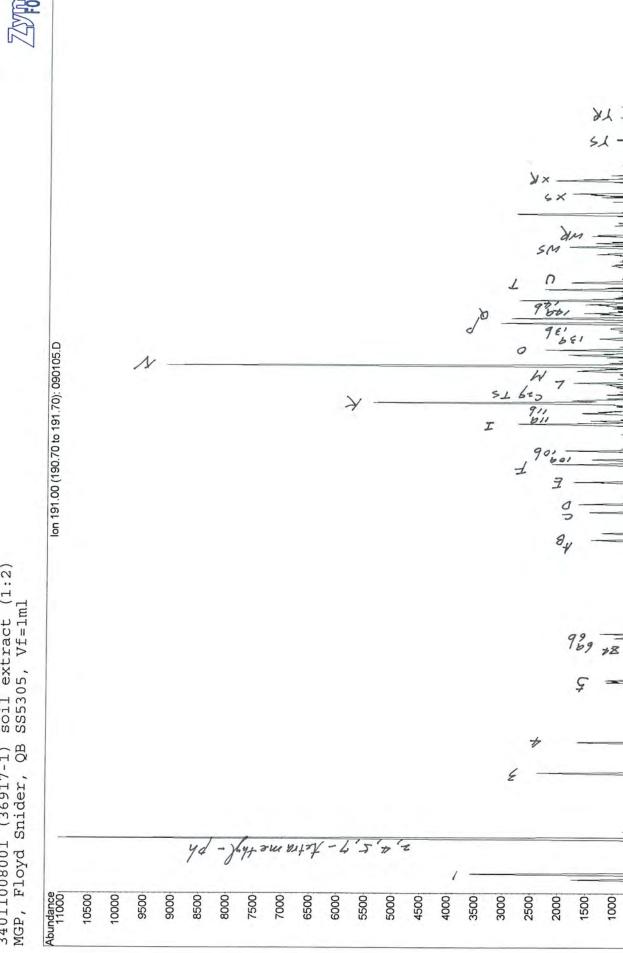
Table



Key for Tricyclic, Tetracyclic, and Pentacyclic Terpanes Identification (m/z 191 mass chromatograms)

Code	Identity	Carbon #		
0	C ₂₀ -Tricyclic Terpane	20		
1	C ₂₁ -Tricyclic Terpane	21		
2 3	C ₂₂ -Tricyclic Terpane	22		
3	C ₂₃ -Tricyclic Terpane	23		
4	C ₂₄ -Tricyclic Terpane	24		
5	C ₂₅ -Tricyclic Terpane	25		
Z4	C24-Tetracyclic Terpane	24		
6a	C ₂₆ -Tricyclic Terpane	26		
6b	C ₂₆ -Tricyclic Terpane	26		
7	C ₂₇ -Tricyclic Terpane	27		
A	C ₂₈ -Tricyclic Terpane #1			
3	C_{28} -Tricyclic Terpane #2	28		
	C_{28} -Tricyclic Terpane #2 C_{29} -Tricyclic Terpane #1	28		
5		29		
Ξ	C ₂₉ -Tricyclic Terpane #2	29		
	18α-22,29,30-Trisnorneohopane (Ts)	27		
3	17α-22,29,30-Trisnorhopane (Tm)	27		
	17ß-22,29-30-Trisnorhopane	27		
H	17α-23,28-Bisnorlupane	28		
10a	C ₃₀ -Tricyclic Terpane #1	30		
10b	C ₃₀ -Tricyclic Terpane #2	30		
Ser 1	17α-28,30-Bisnorhopane	28		
11a	C ₃₁ -Tricyclic Terpane #1	31		
	17α-25-Norhopane	29		
1b	C ₃₁ -Tricyclic Terpane #2	31		
<	17α,21β-30-Norhopane	29		
C ₂₉ Ts	18α-30-Norneohopane	29		
230*	17α-Diahopane	30		
	17β-21α-30-Normoretane	29		
la	18α-Oleanane	30		
٨b	18ß-Oleanane	30		
1	17α ,21ß-Hopane	30		
)	17β,21α-Moretane	30		
3a	C ₃₃ -Tricyclic Terpane #1			
3b	C ₃₃ -Tricyclic Terpane #2	33		
	22S-17α,21β-30-Homohopane	33		
1		31		
	22R-17α,21β-30-Homohopane	31		
4a	Gammacerane	30		
44	C ₃₄ -Tricyclic Terpane #1	34		
	17β,21α-Homomoretane	31		
4b	C ₃₄ -Tricyclic Terpane #2	34		
	22S-17α,21β-30-Bishomohopane	32		
	22R-17α,21ß-30-Bishomohopane	32		
5a	C ₃₅ -Tricyclic Terpane #1	35		
ōb	C ₃₅ -Tricyclic Terpane #2	35		
	17ß,21α-C ₃₂ -Bishomomoretane	32		
S	22S-17α,21β-30,31,32-Trishomohopane	33		
R	22R-17α,21β-30,31,32-Trishomohopane	33		
ba -	C ₃₆ -Tricyclic Terpane #1	36		
6b	C ₃₆ -Tricyclic Terpane #2	36		
5	22S-17α,21ß-30,31,32,33-Tetrahomohopane	34		
2	22R-17a,21ß-30,31,32,33-Tetrahomohopane	34		
6	22S-17α,21β-30,31,32,33,34-Pentahomohopane	35		
2	22R-17α,21β-30,31,32,33,34-Pentahomohopane	35		

soil extract (1:2) SS5305, Vf=lml 34011008001 (36917-1) MGP, Floyd Snider, QB



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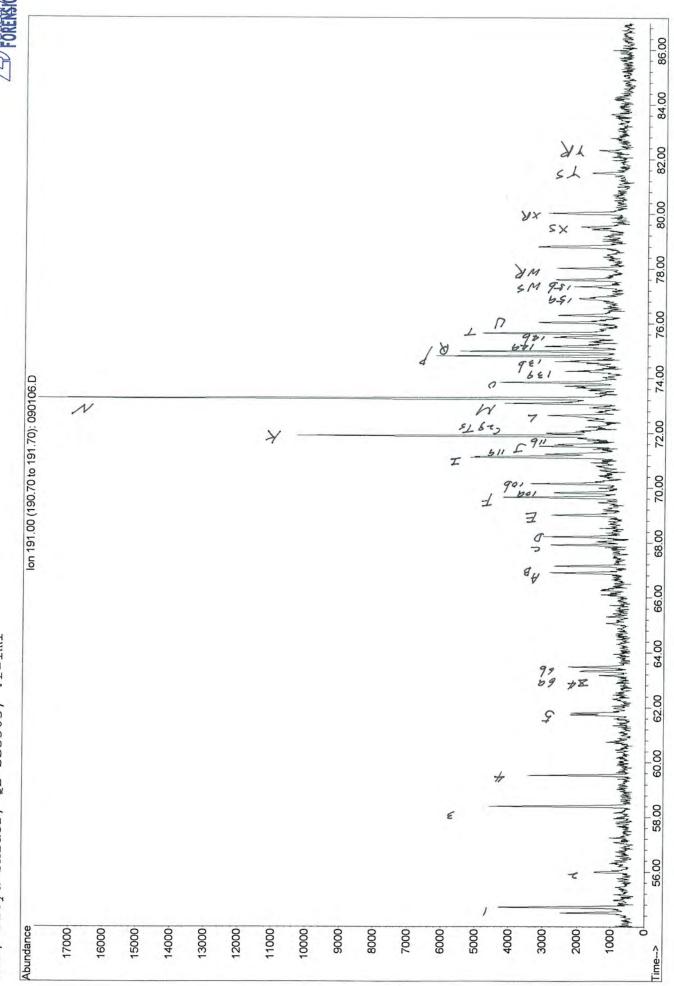
3

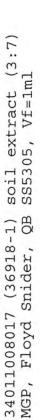
and have been a product of the test of test of

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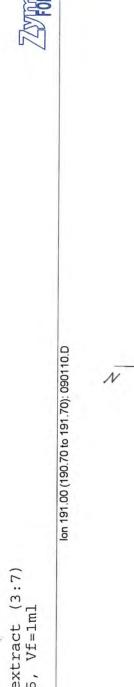
51

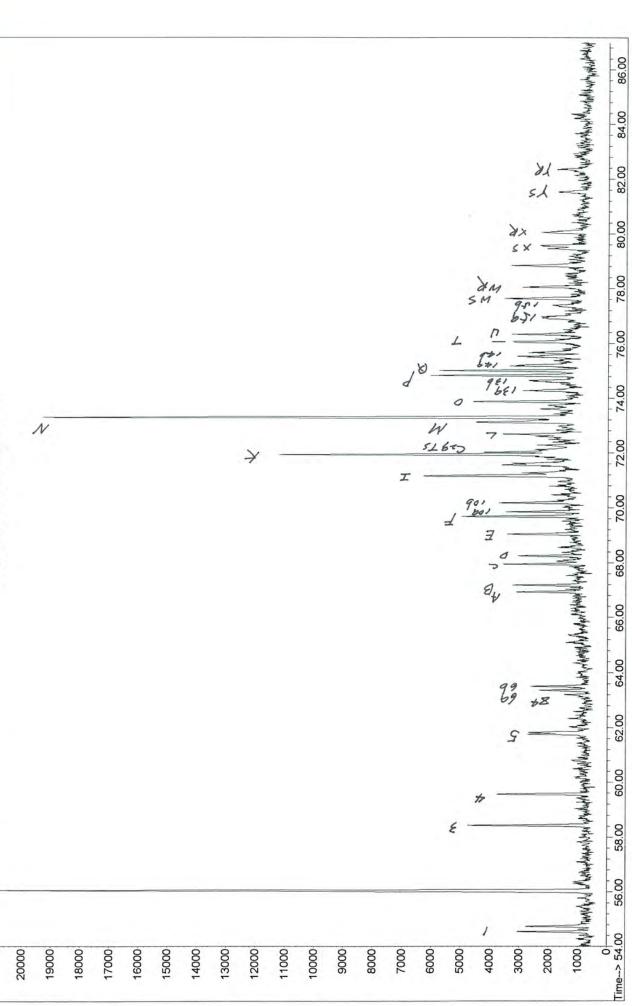
34011008011 (36917-11) soil extract (1:2) MGP, Floyd Snider, QB SS5305, Vf=1ml





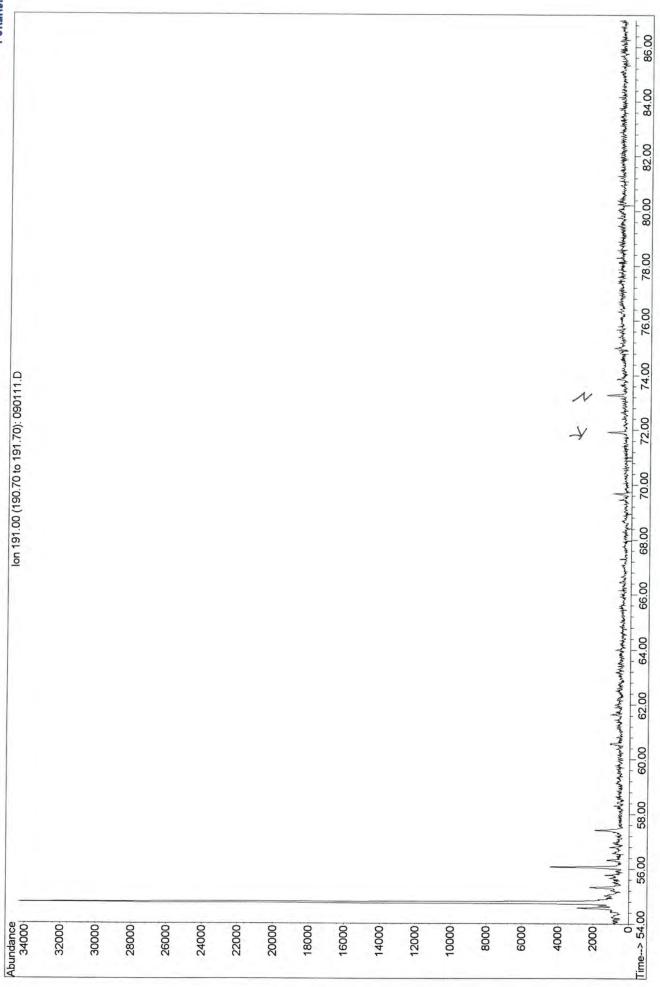
Abundance



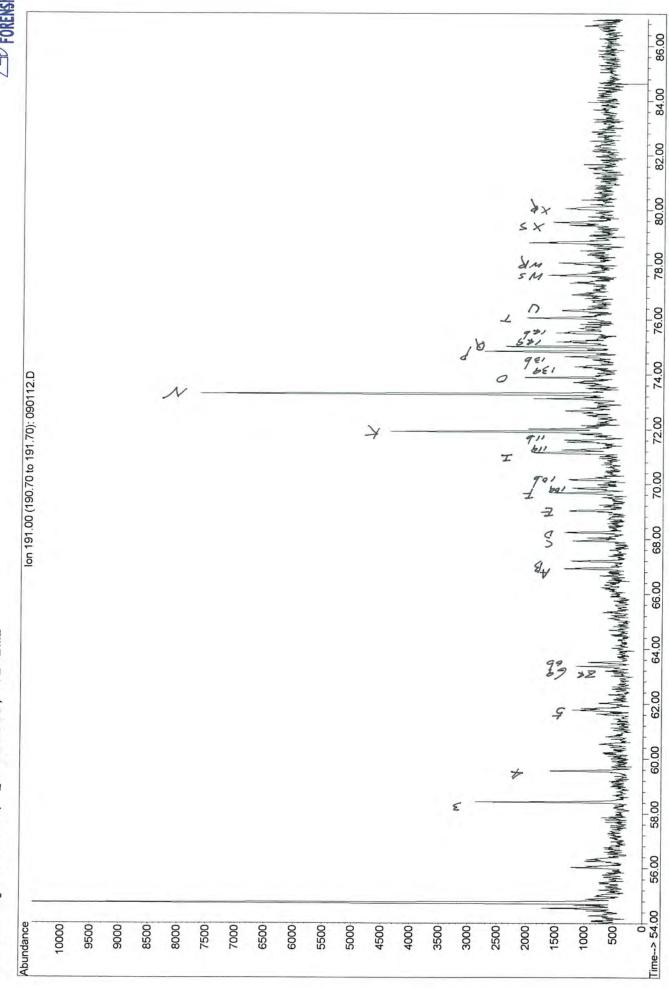


34011008020 (36918-4) soil extract (1:1.2) MGP, Floyd Snider, QB SS5305, Vf=2ml



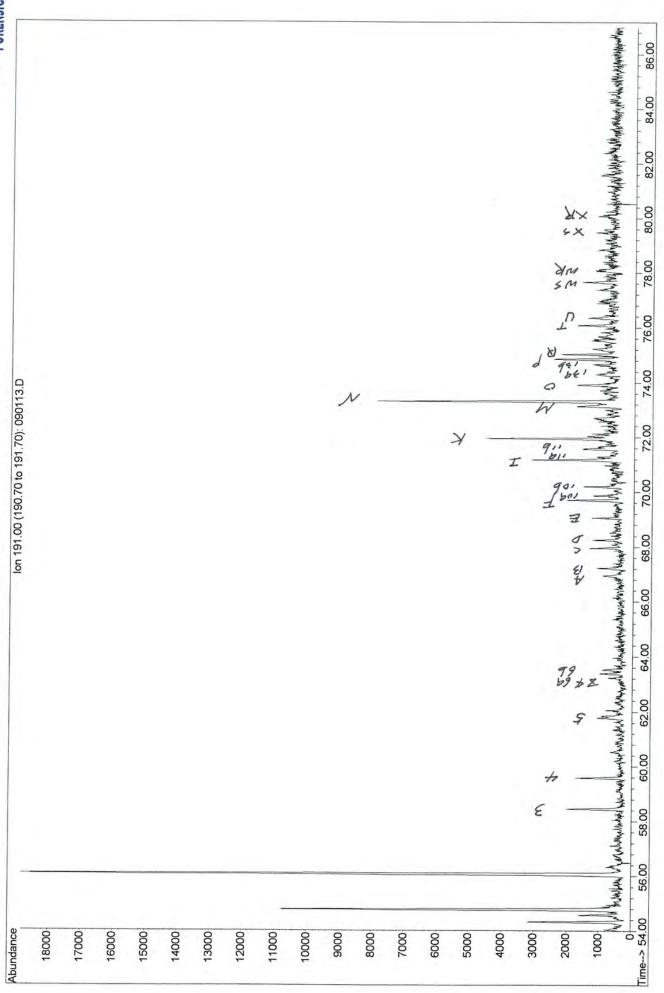


34011008022 (36918-6) soil extract (3:8) MGP, Floyd Snider, QB SS5305, Vf=2ml



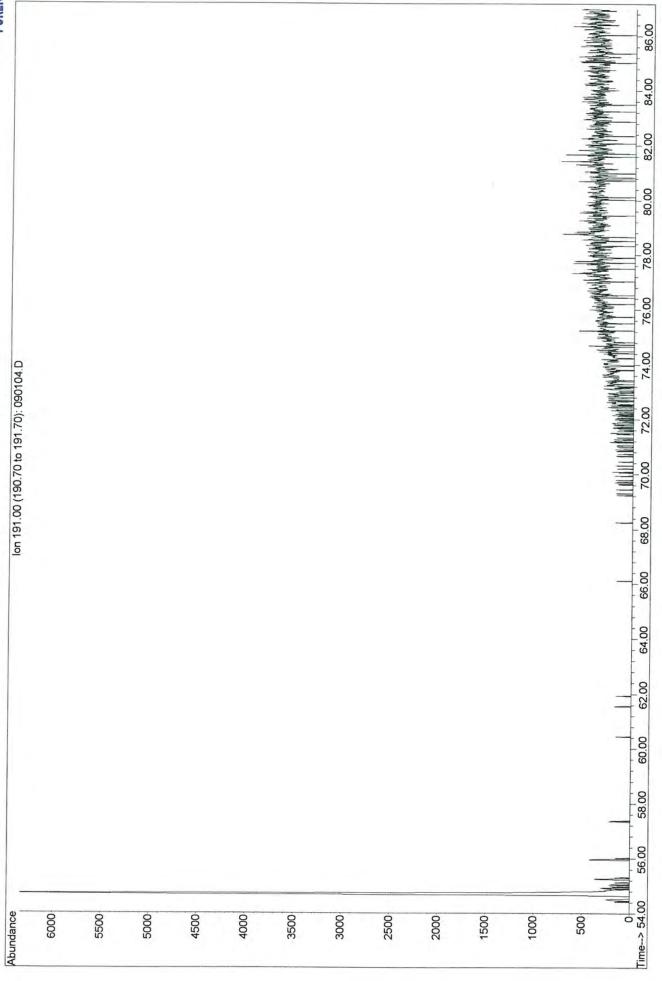
34011008028 (36918-12) soil extract (1:1)) MGP, Floyd Snider, QB SS5305, Vf=2ml





34011008030 (36918-14) soil extract MGP, Floyd Snider, QB SS5305





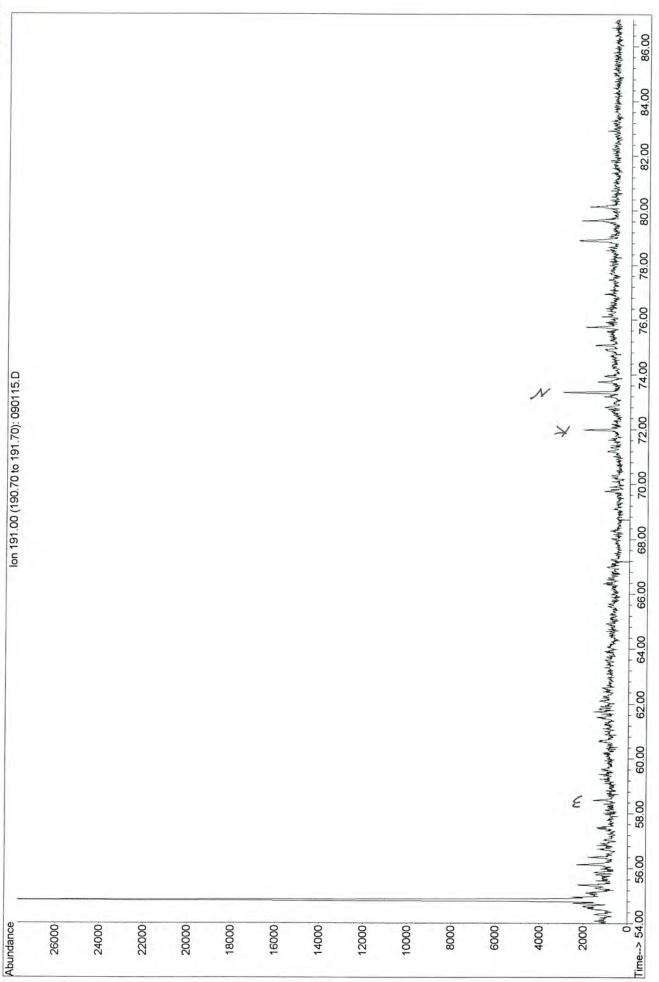
34011008032 (36918-16) soil ext (3.5:6.5) MGP, Floyd Snider, QB SS5305, Vf=4ml



		the second
		WWWWWWWWWWWWWWWWWWWWWWWWWWWWWWWWWWWWWW

34011008035 (36918-19) soil ext (4:6) MGP, Floyd Snider, QB SS5305, Vf=1ml





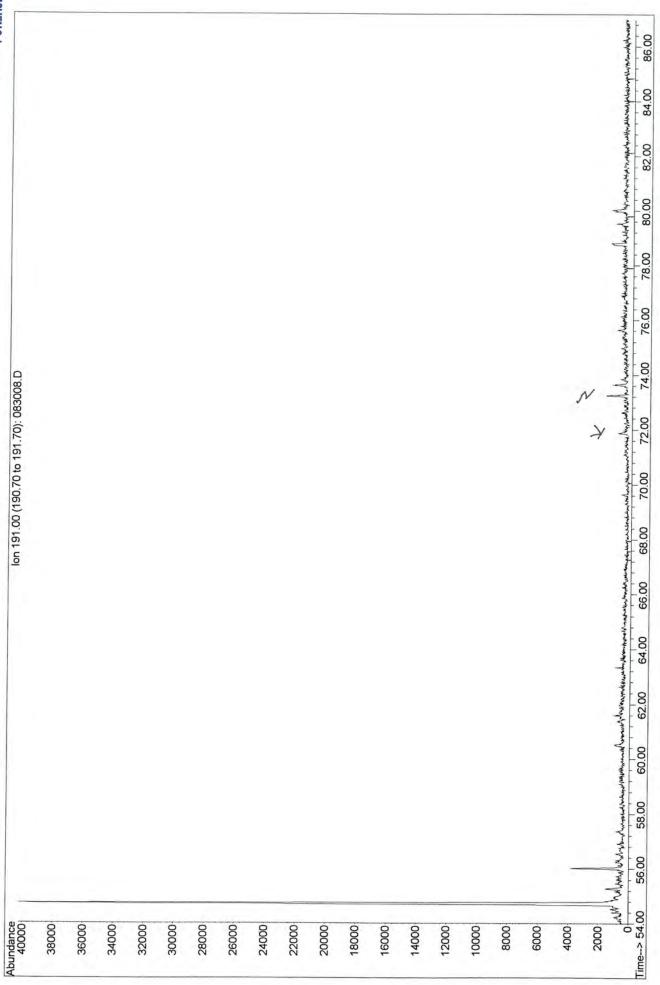
34011008040 (36919-4) soil extract (1:2) MGP, Floyd Snider, QB SS5318



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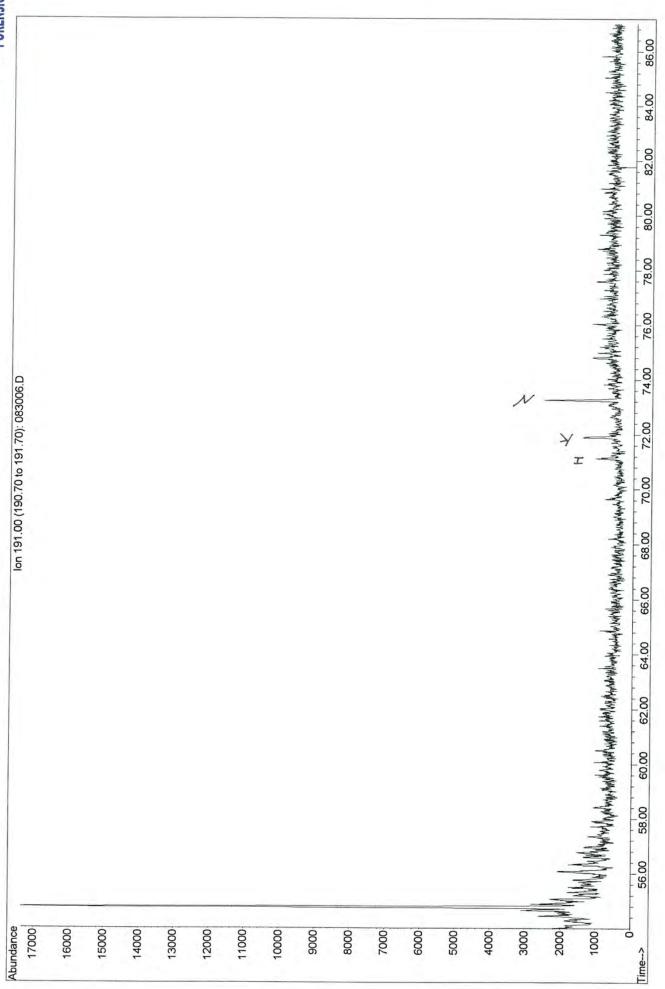
34011008043 (36919-7) soil extract (1:2) MGP, Floyd Snider, QB SS5318, Vf=3ml





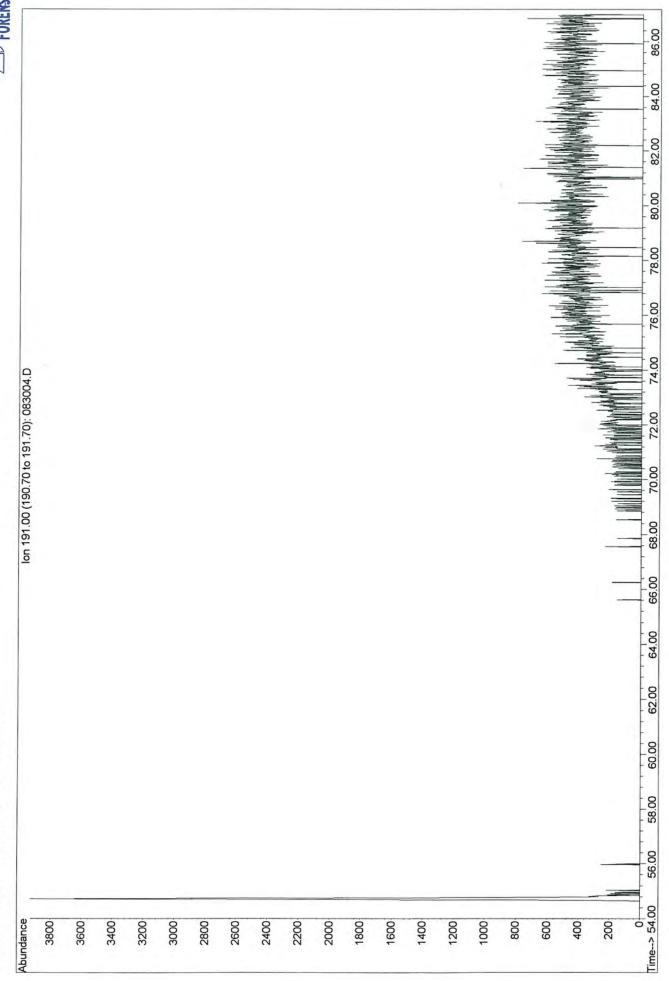
34011008050 (36920-1) soil extract (1:7) MGP, Floyd Snider, QB SS5318, Vf=3ml

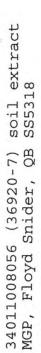




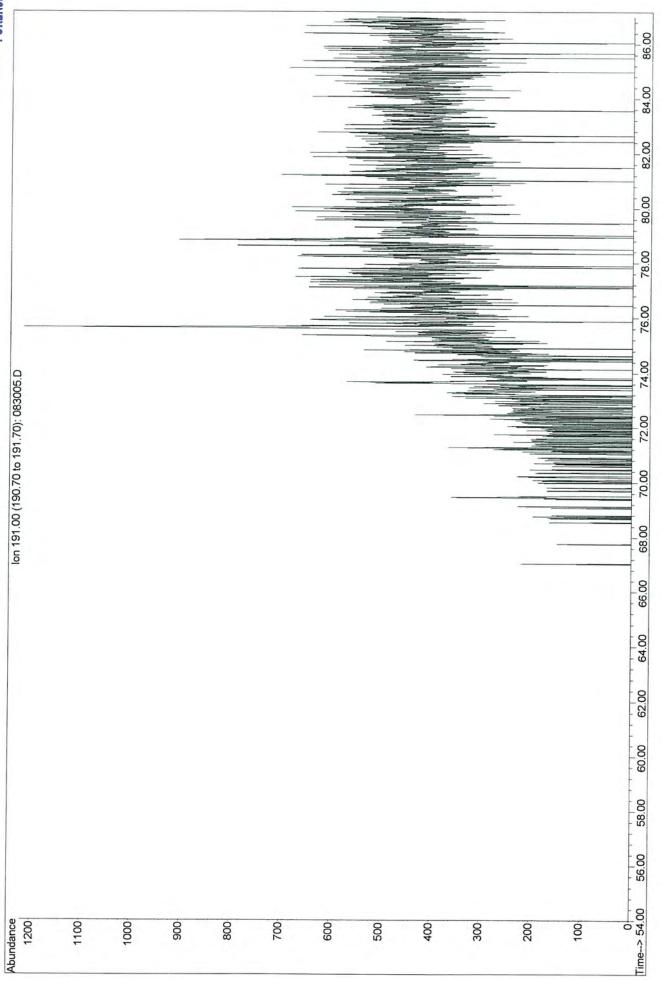
34011008054 (36920-5) soil extract MGP, Floyd Snider, QB SS5318





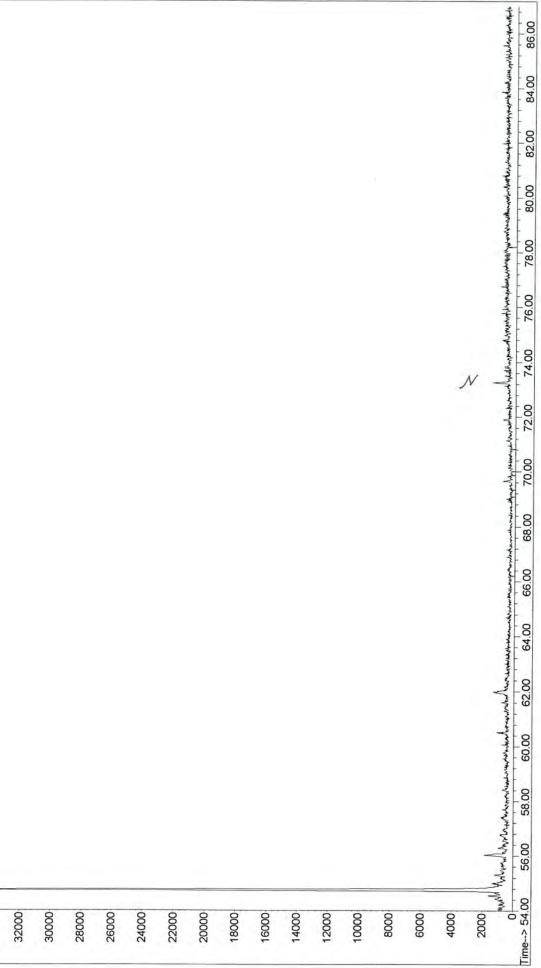






DW-4-FP (36921-1) Product x50 Gas Works Park, Flovd

Abundance Ian 191.00 (190.70 to 191.70): 090305.D 38000 38000 34000 32000 32000 28000 28000 28000



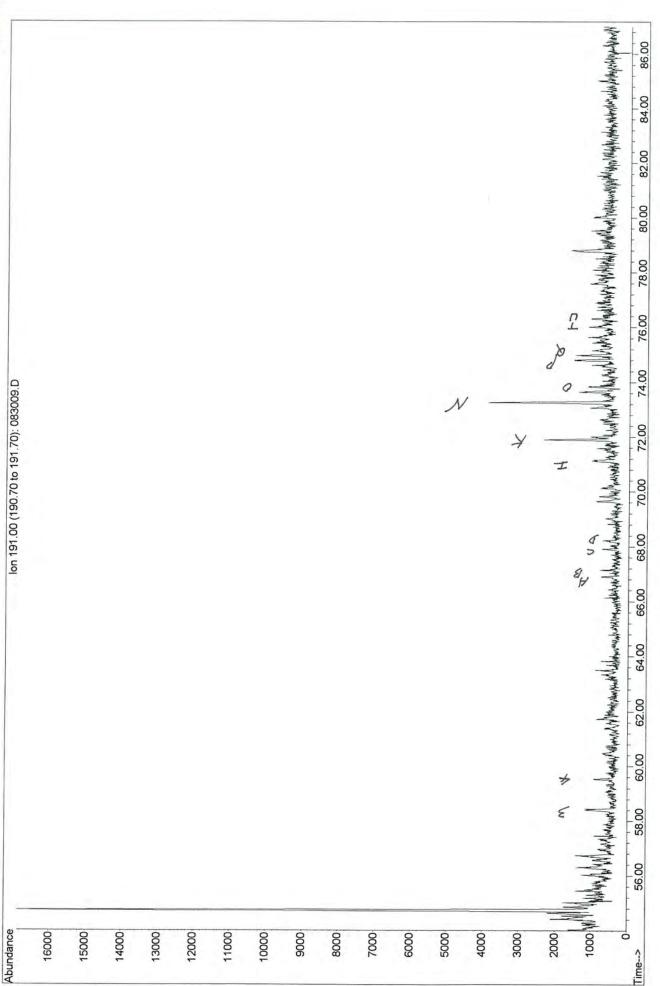


DW-5-FP (36921-2) Product x50 Gas Works Park, Floyd



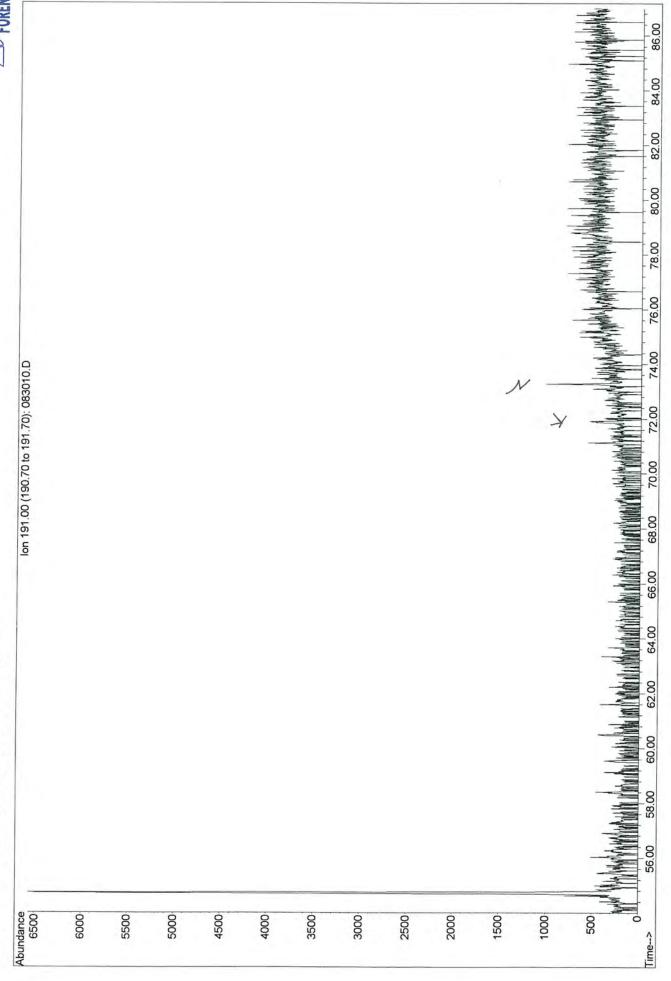


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1-1G (36922-3) soil extract (1:12) MGP, Floyd Snider, QB SS5318, Vf=3ml





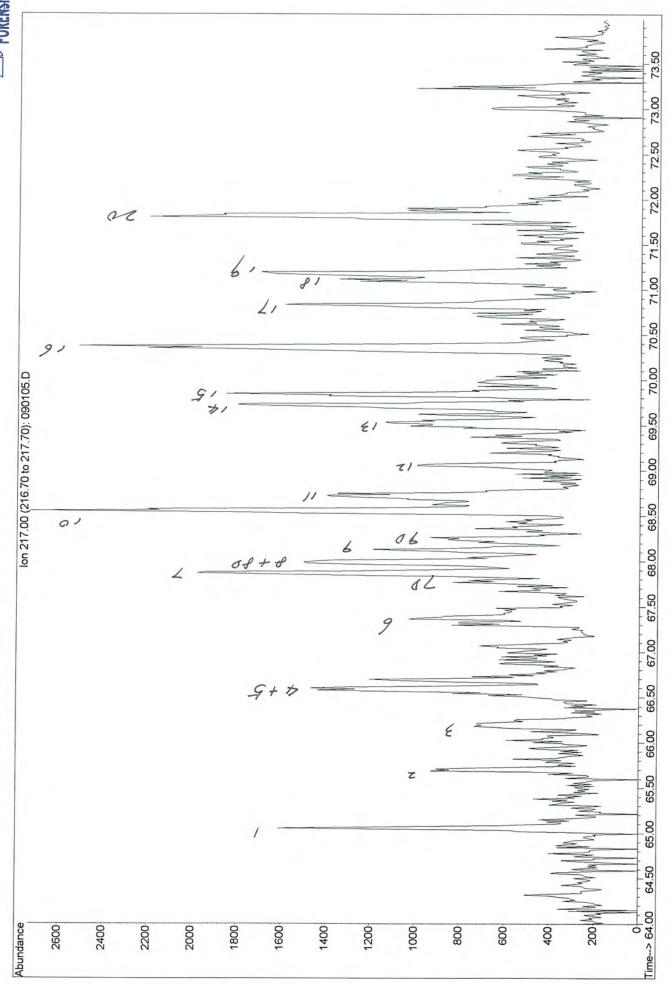
Table

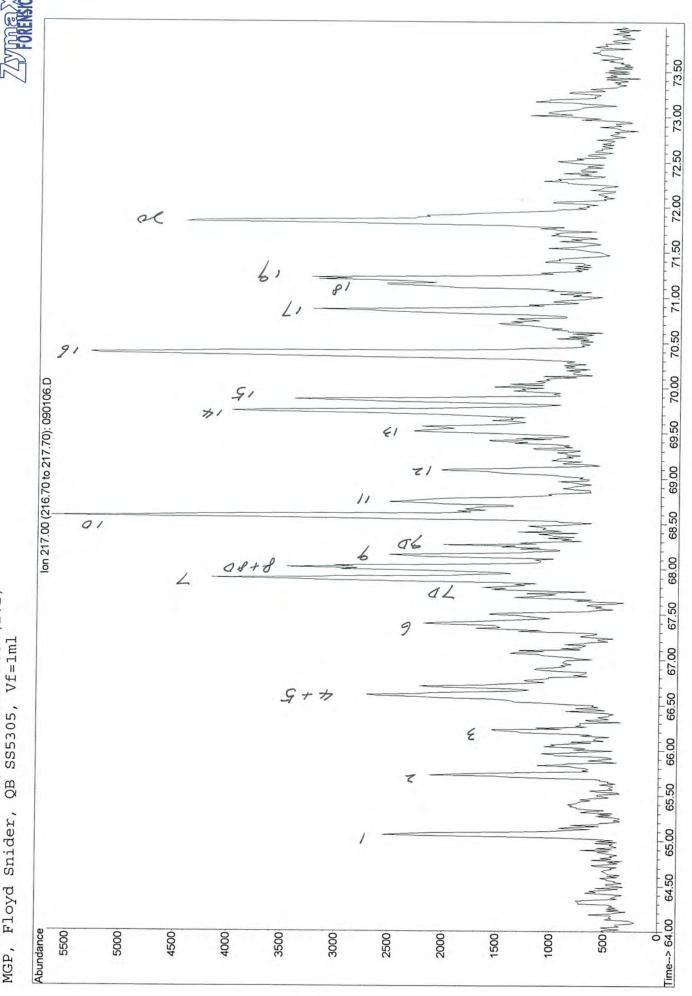


Key for Steranes Identification (m/z 217 Mass Chromatogram)

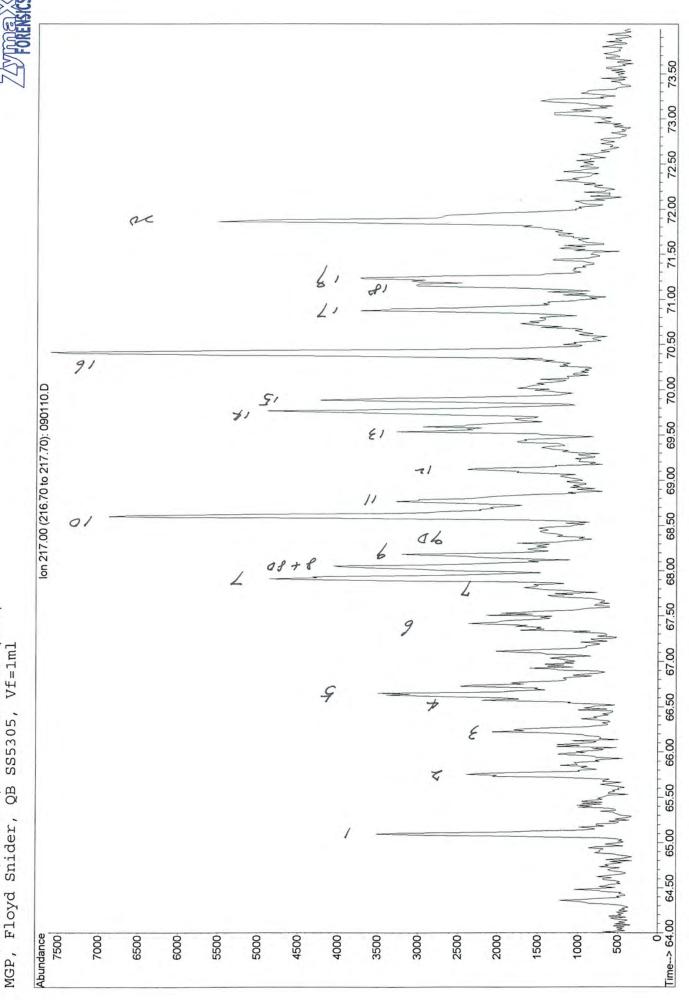
Code	Identity	Carbon #			
1	13ß,17α-diacholestane (20S)	27			
2	13ß,17α-diacholestane (20R)	27			
3	13α,17ß-diacholestane (20S)	27			
4	13α,17β-diacholestane (20R)	27			
5	24-methyl-13ß,17α-diacholestane (20S)	28			
6	24-methyl-13ß,17α-diacholestane (20R)	28			
7D	24-methyl-13α,17β-diacholestane (20S)	28			
7	14α,17α-cholestane (20S)	27			
8D	24-ethyl-13ß,17α-diacholestane (20S)	29			
8	14ß,17ß-cholestane (20R)	27			
9	14ß,17ß-cholestane (20S)	27			
9D	24-methyl-13α,17β-diacholestane (20R)	28			
10	14α,17α-cholestane (20R)	27			
11	24-ethyl-13ß,17α-diacholestane (20R)	29			
12	24-ethyl-13α,17β-diacholestane (20S)	29			
13	24-methyl-14α,17α-cholestane (20S)	28			
4D	24-ethyl-13α,17β-diacholestane (20R)	29			
4	24-methyl-14ß,17ß-cholestane (20R)	28			
5	24-methyl-14ß,17ß-cholestane (20S)	28			
6	24-methyl-14α,17α-cholestane (20R)	28			
7	24-ethyl-14α-cholestane (20S)	29			
8	24-ethyl-14ß,17ß-cholestane (20R)	29			
9	24-ethyl-14ß,17ß-cholestane (20S)	29			
0	24-ethyl-14α,17α-cholestane (20R)	29			
1A	24-n-Propylcholestane (20S)	30			
1B	4-methyl-24-ethylcholestane (20S)	30			
2A	4α -methyl-24-ethyl-14 β ,17 β -cholestane(20S)	30			
2B	24-n-propyl-14β,17β-cholestane (20S)	30			
3A	4α -methyl-24-ethyl-14 β ,17 β -cholestane(20R)	30			
BB	24-n-propyl-14β,17β-cholestane (20R)	30			
1A	4α-methyl-24-ethylcholestane(20R)	30			
B	24-n-propylcholestane (20R)	30			







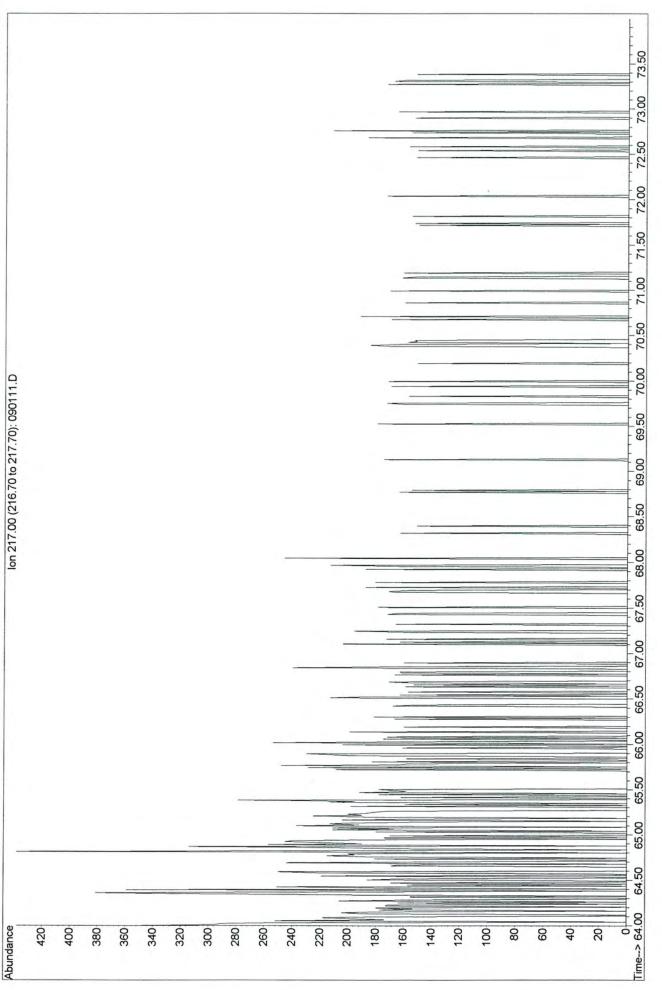
34011008011 (36917-11) soil extract (1:2) MGP, Floyd Snider, QB SS5305, Vf=1ml

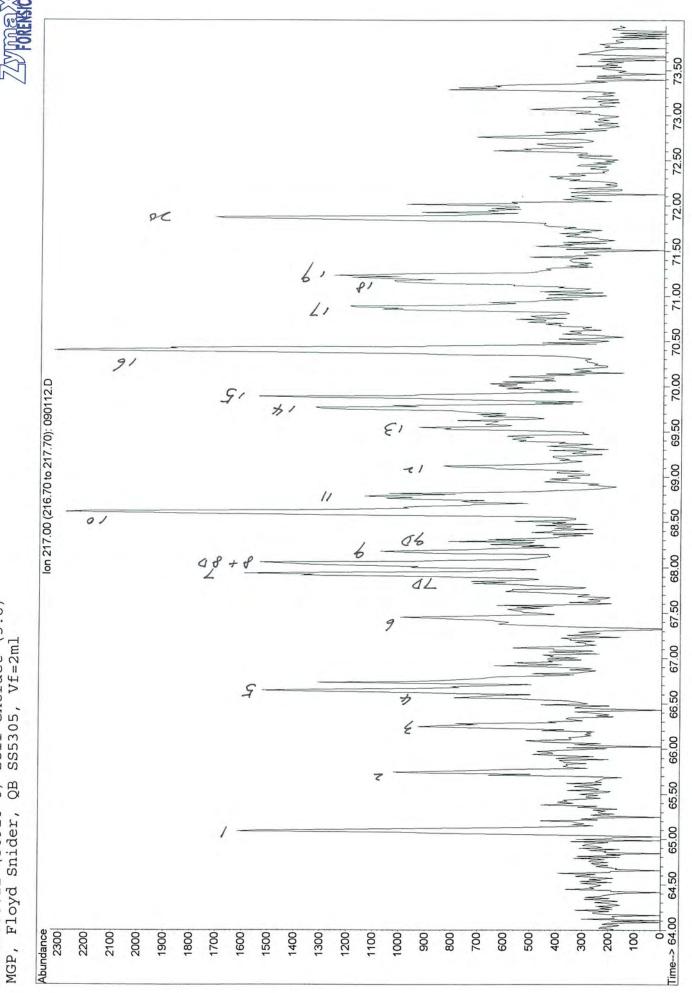


soil extract (3:7)
SS5305, Vf=lml 34011008017 (36918-1) MGP, Floyd Snider, QB

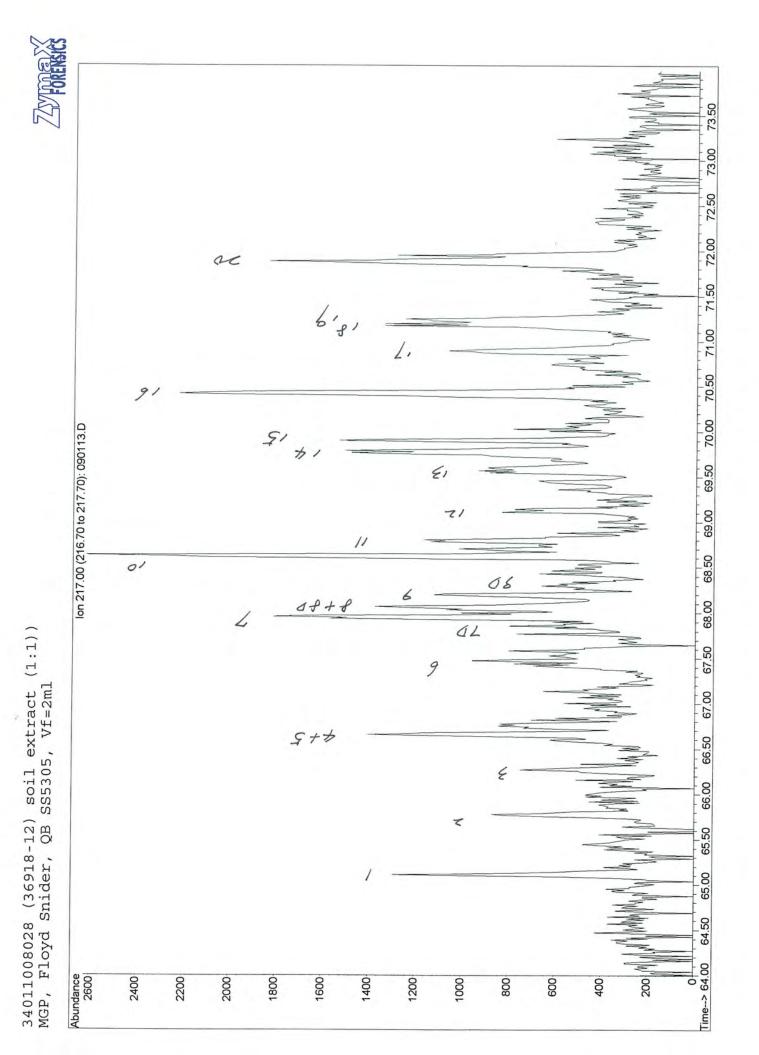
34011008020 (36918-4) soil extract (1:1.2) MGP, Floyd Snider, QB SS5305, Vf=2ml





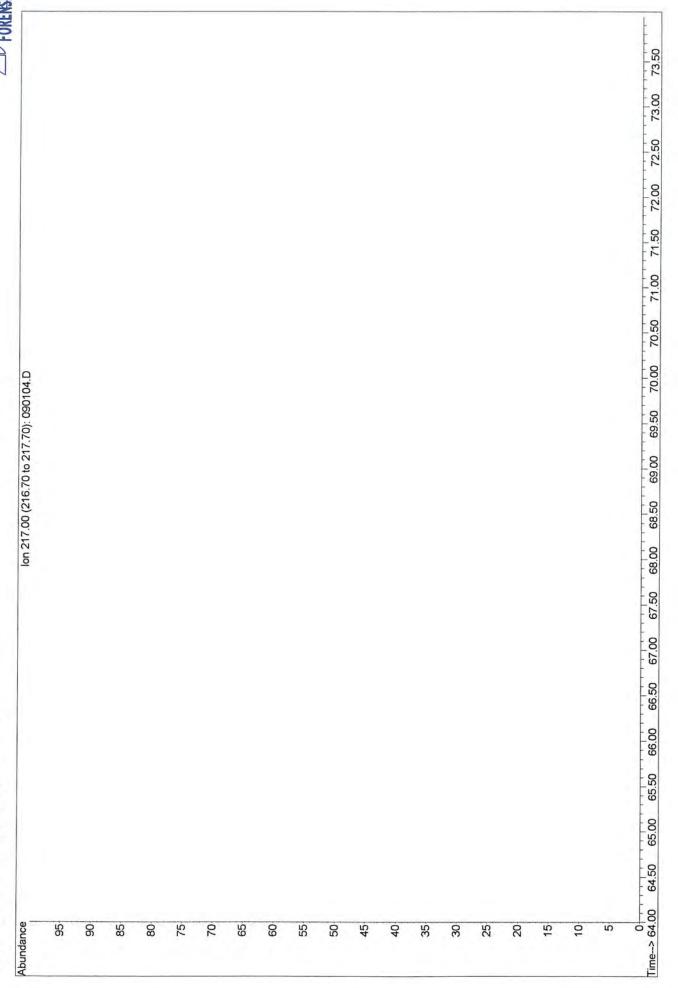


soil extract (3:8)
SS5305, Vf=2ml 34011008022 (36918-6) MGP, Floyd Snider, QB



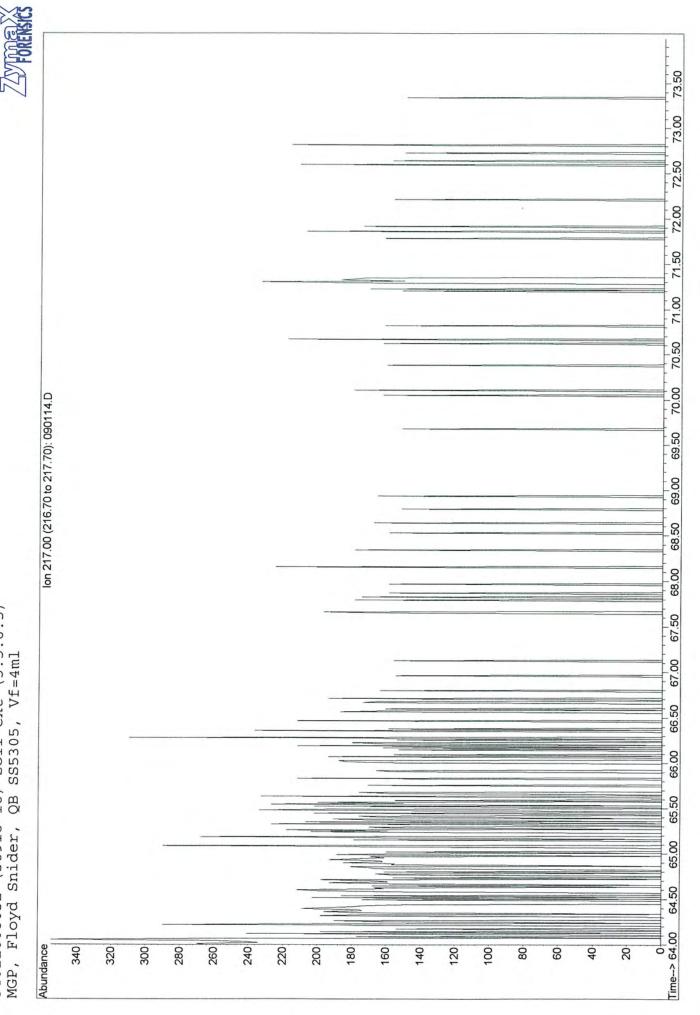
34011008030 (36918-14) soil extract MGP, Floyd Snider, QB SS5305



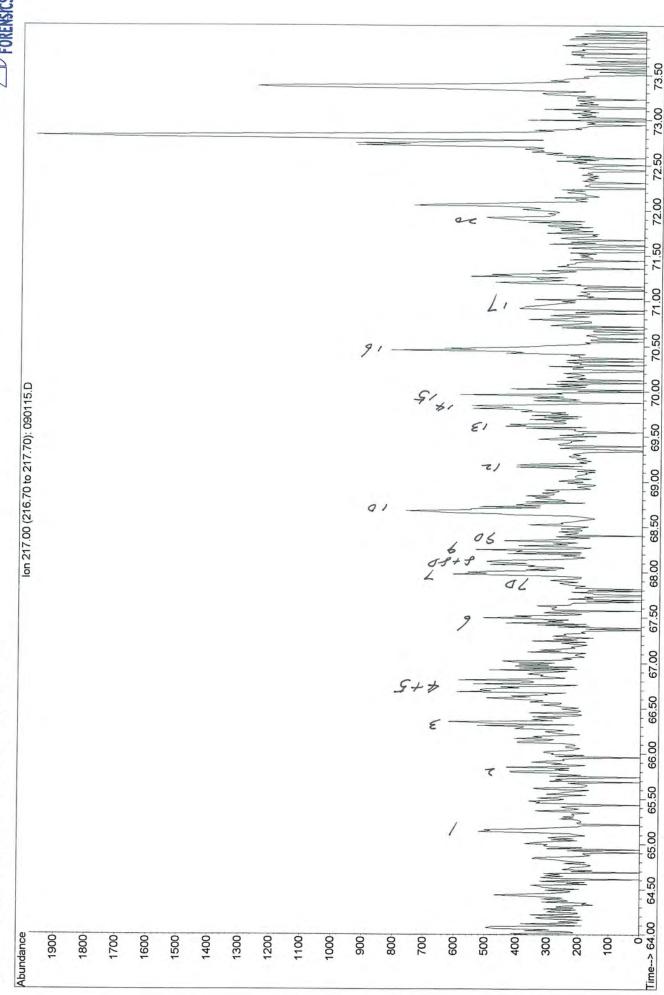


34011008032 (36918-16) soil ext (3.5:6.5) MGP, Floyd Snider, QB SS5305, Vf=4ml





34011008035 (36918-19) soil ext (4:6) MGP, Floyd Snider, QB SS5305, Vf=1ml



34011008040 (36919-4) soil extract (1:2) MGP, Floyd Snider, QB SS5318

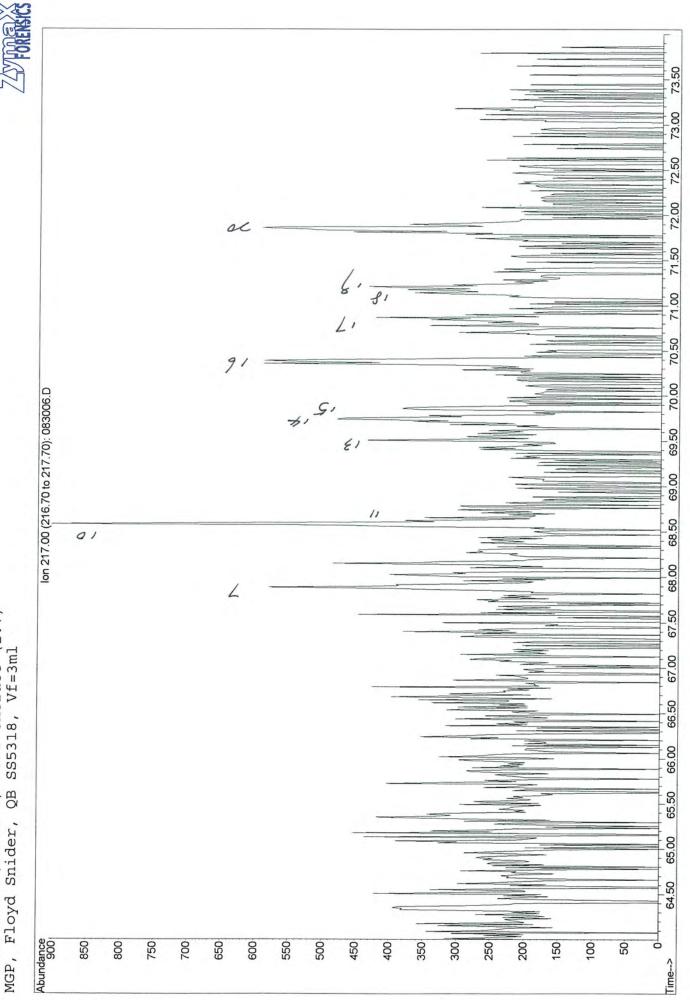
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34011008043 (36919-7) soil extract (1:2) MGP, Floyd Snider, QB SS5318, Vf=3ml

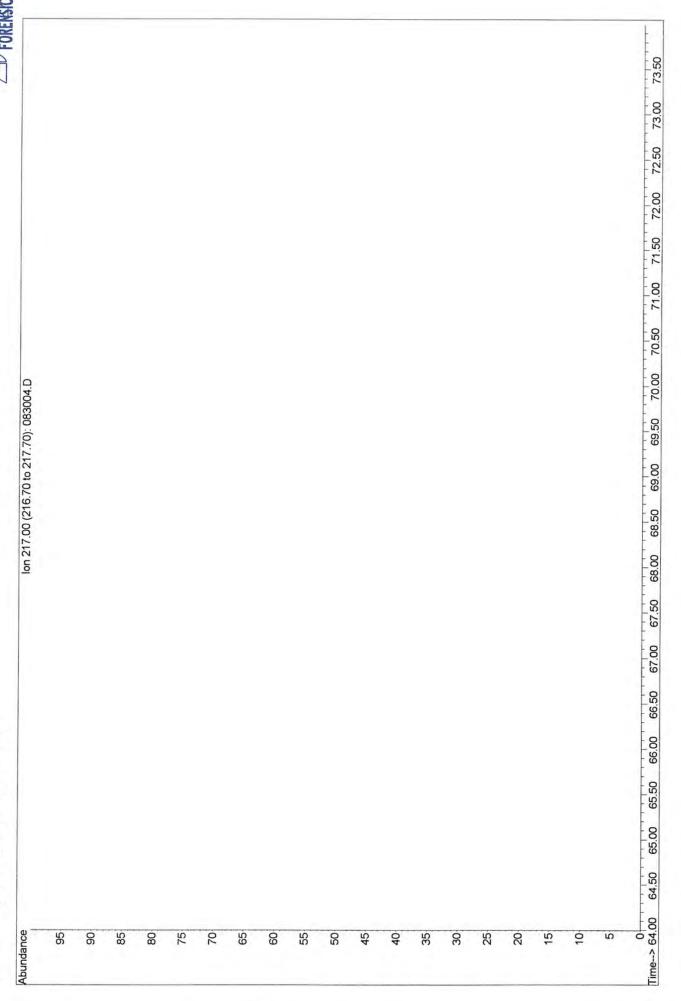
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7.70): 08									-			
70 to 21								-				
0 (216.												
lon 217.00 (216.70 to 217.70): 083008.D												
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Abundance	600	550	500	450	400	350	300	250	200	150	100	2 2

soil extract (1:7) SS5318, Vf=3ml 34011008050 (36920-1) MGP, Floyd Snider, QB



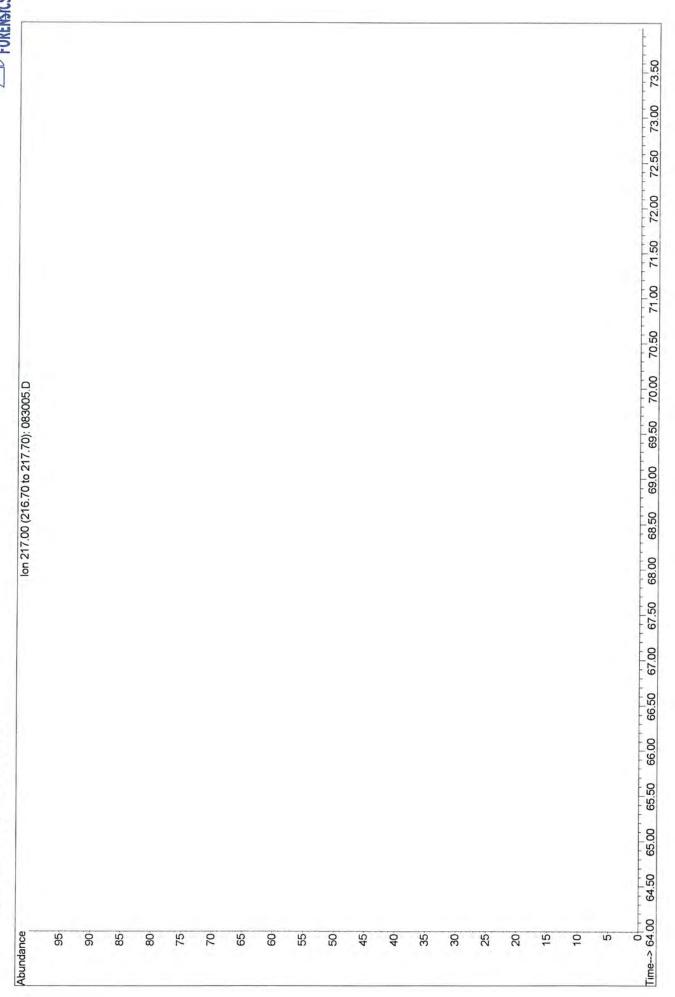
34011008054 (36920-5) soil extract MGP, Floyd Snider, QB SS5318



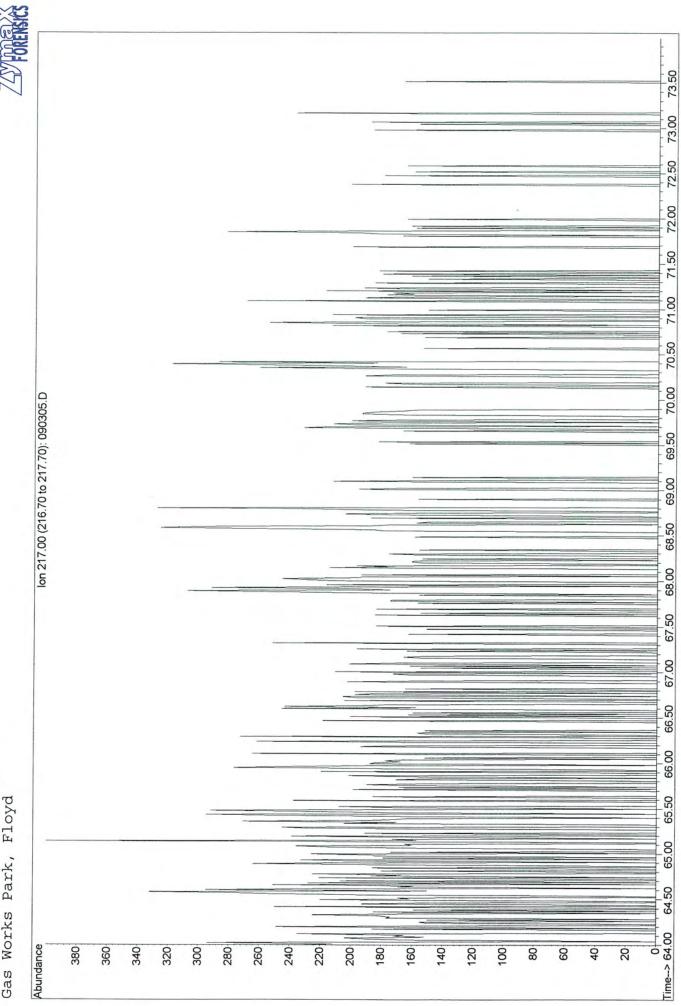


34011008056 (36920-7) soil extract MGP, Floyd Snider, QB SS5318

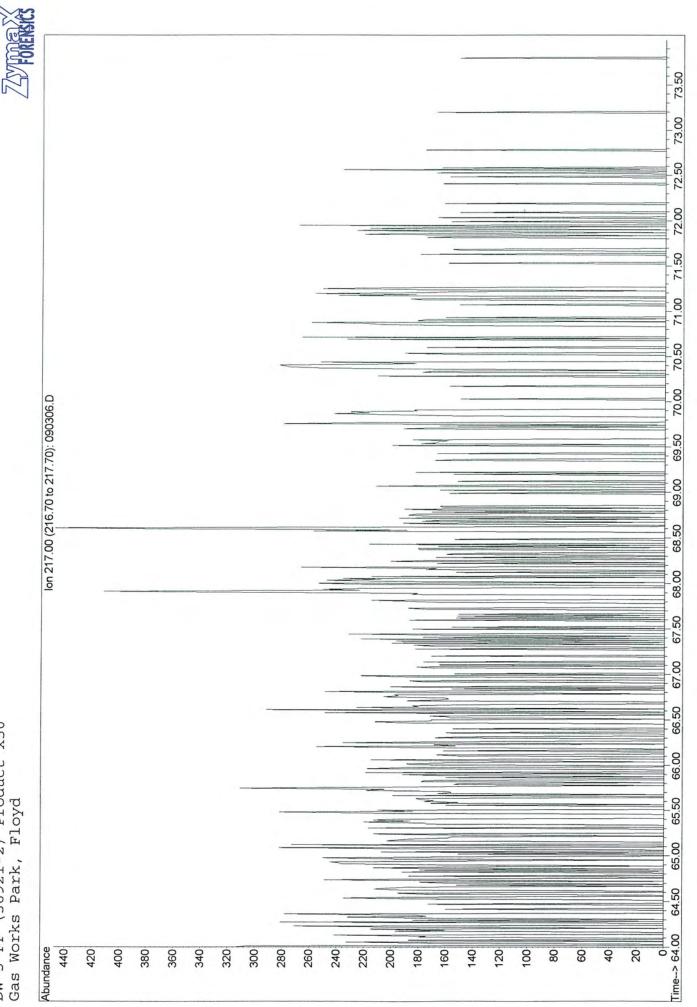


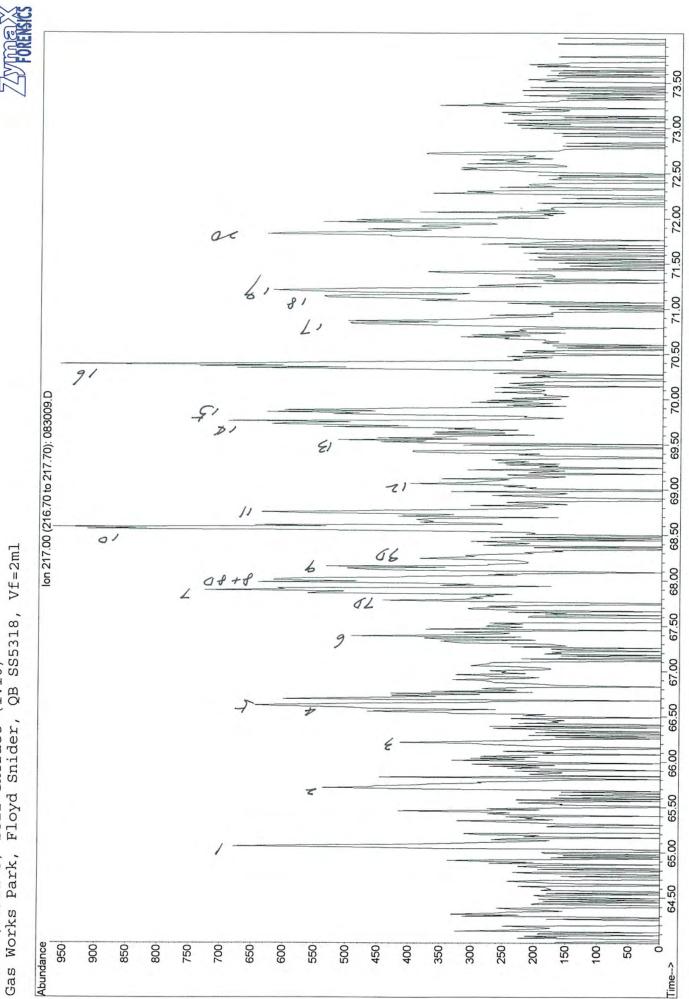






DW-5-FP (36921-2) Product x50 Gas Works Park, Floyd





soil extract (1:10)
Floyd Snider, QB SS5318, Vf=2ml 3-1-G (36921-3) Gas Works Park,

1-1G (36922-3) soil extract (1:12) MGP, Floyd Snider, QB SS5318, Vf=3ml



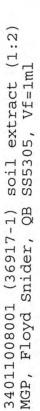
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83010.D			_	_			-															
217.70): 0																						
lon 217.00 (216.70 to 217.70): 083010.D						_																
on 217.00																						
							-				_				1.00							
						_						_										
						_																
						1										-						
ADUNDANCE	230	220	210	200	190	180	170	160	150	140	130	120	110	100	90	80	70	60	20	40.	30.30	

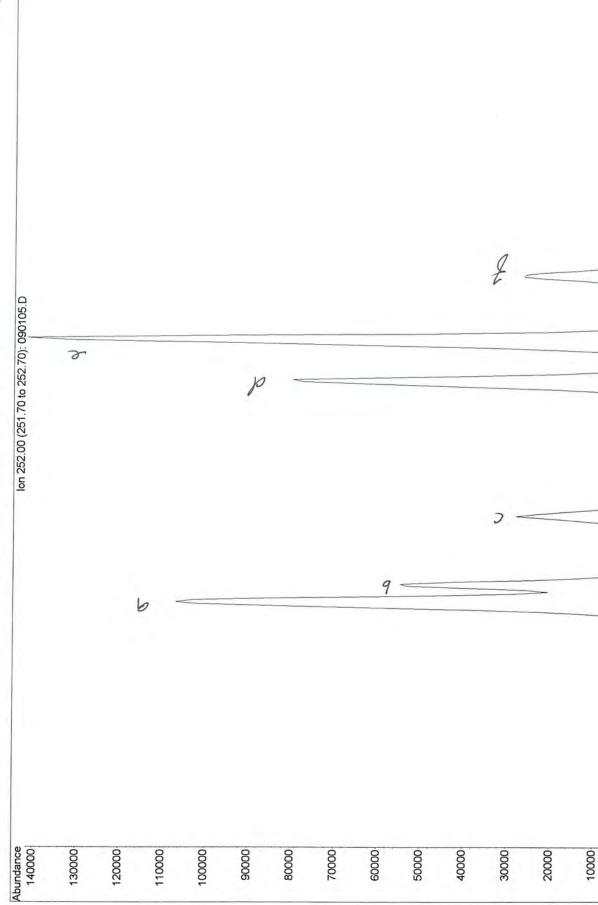


Table

Key for Identification for Six Pyrogenic PAH (m/z 252)

Peak No.	Identity
а	Benzo(B)fluoranthene
b	Benzo(K)fluoranthene
с	Benzo(A)fluoranthene
d	Benzo(E)pyrene
е	Benzo(A)pyrene
f	Perylene

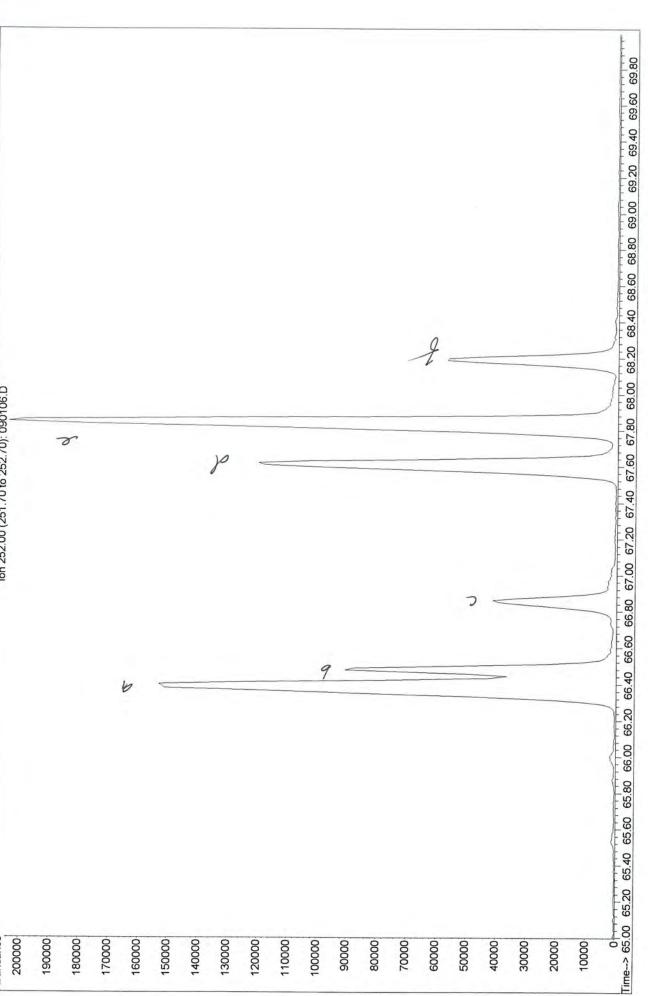


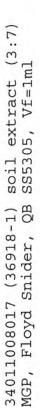


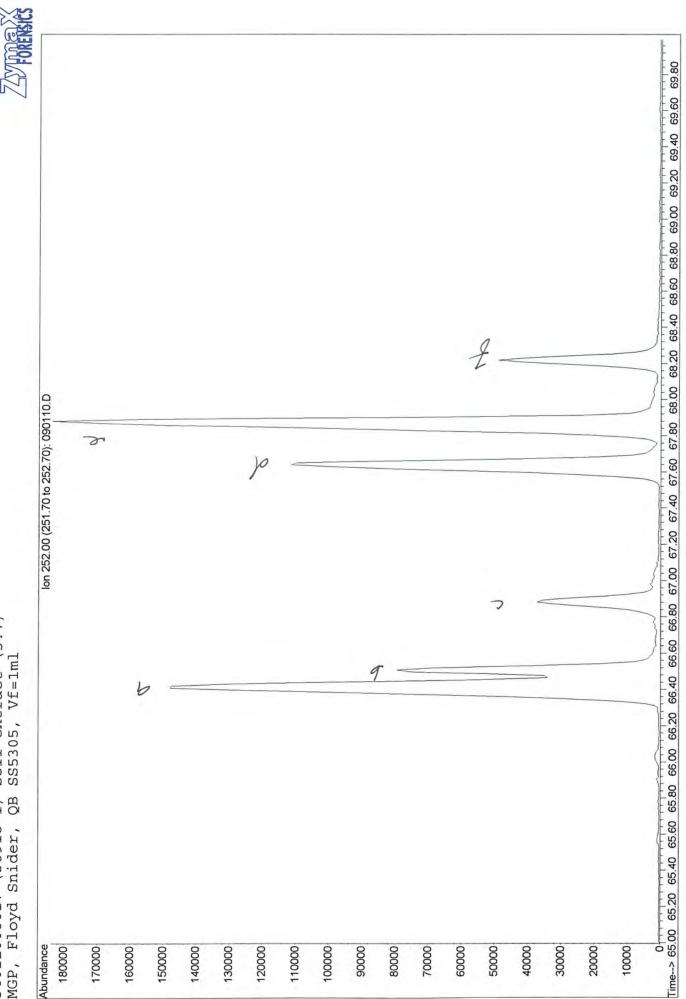
0 Time--> 65.00 65.20 65.40 65.60 65.80 66.00 66.20 66.40 66.60 67.00 67.20 67.40 67.60 67.80 68.00 68.20 68.40 68.60 68.80 69.00 69.20 69.40 69.60 69.60

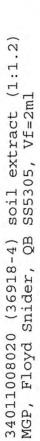


PORENEICS

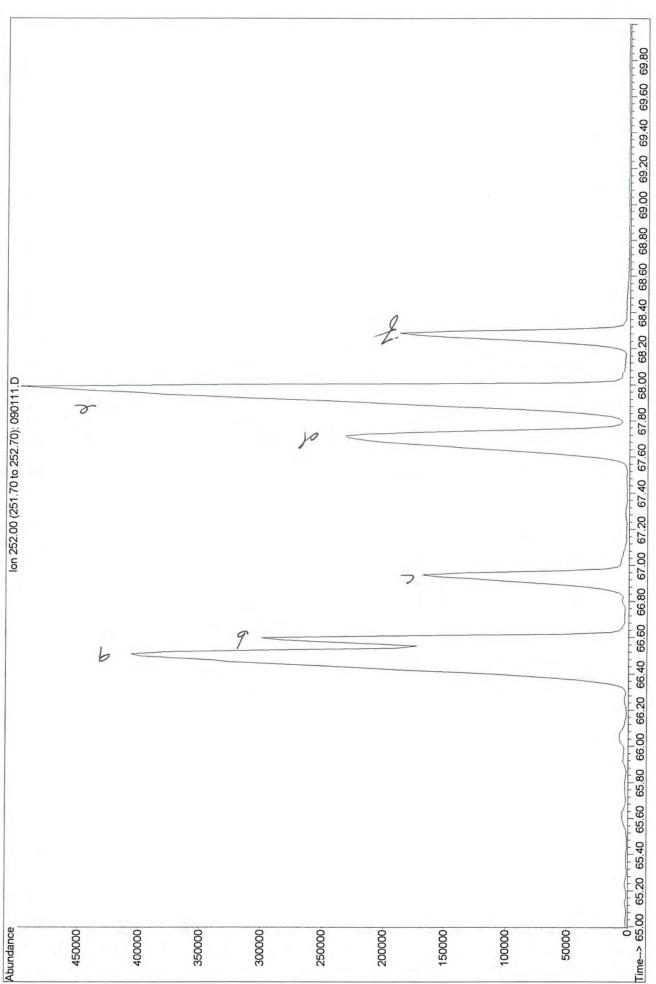


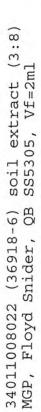






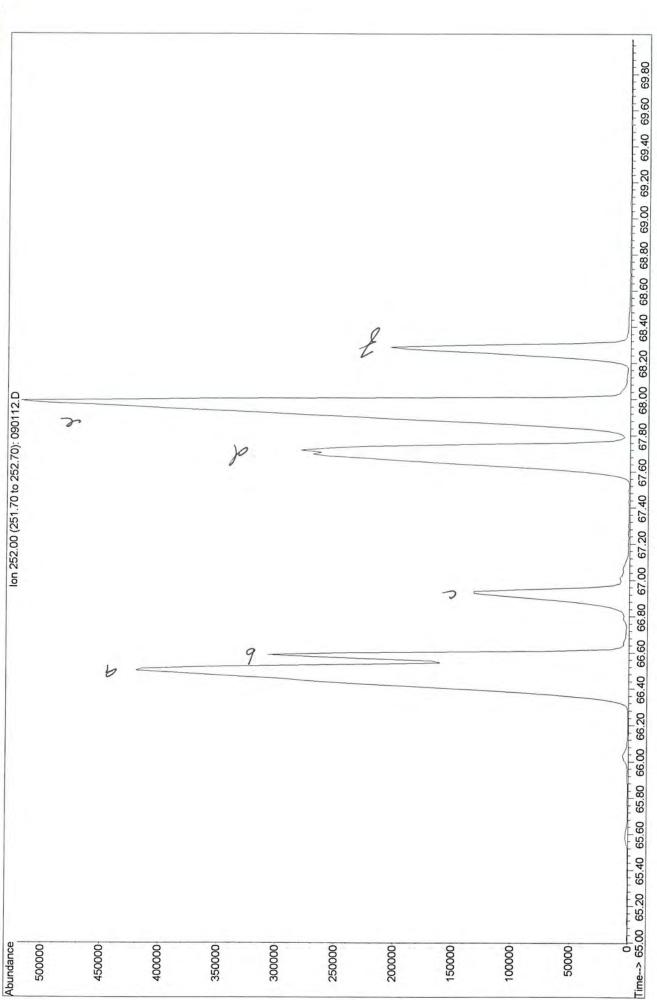




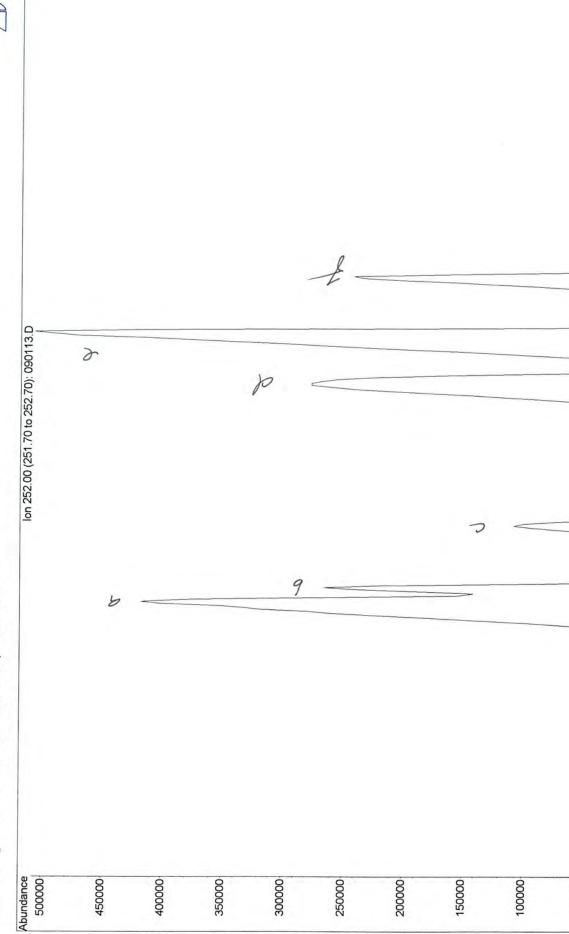




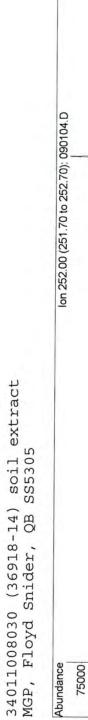
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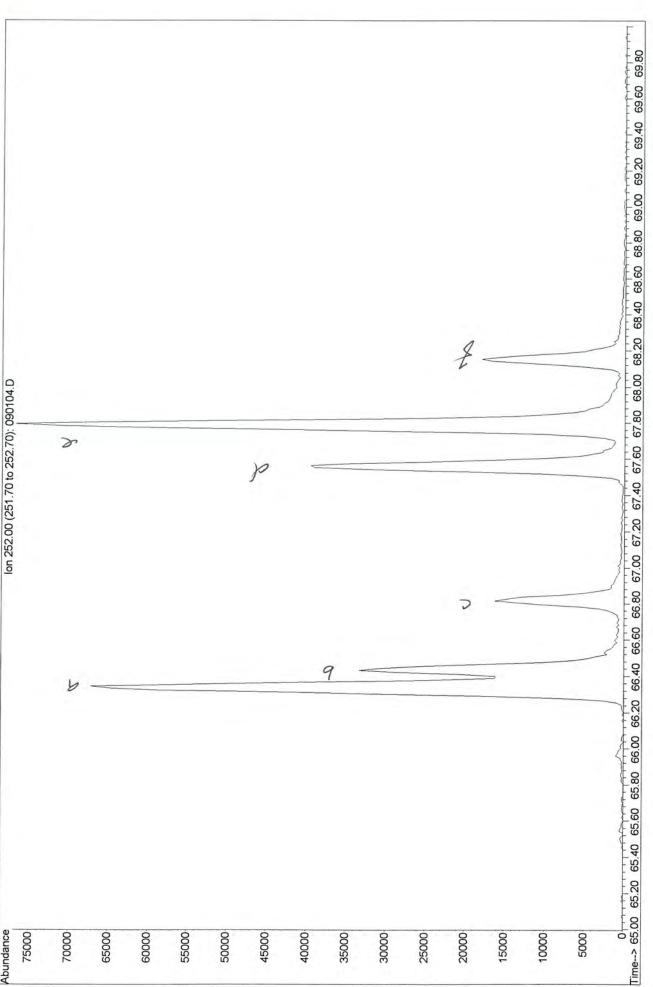




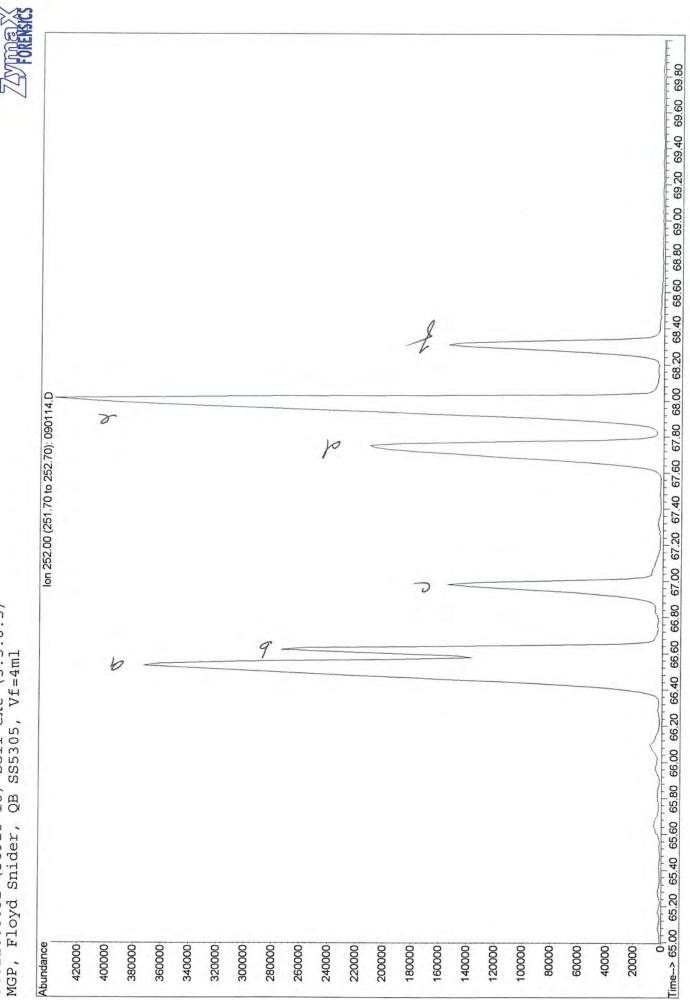


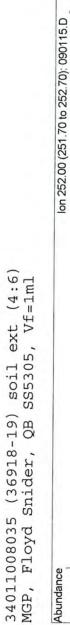
Time--> 65.00 65.20 65.40 65.60 65.80 66.00 66.20 66.40 66.60 67.00 67.20 67.40 67.60 67.80 68.00 68.20 68.40 68.60 68.80 69.00 69.20 69.40 69.60 69.80

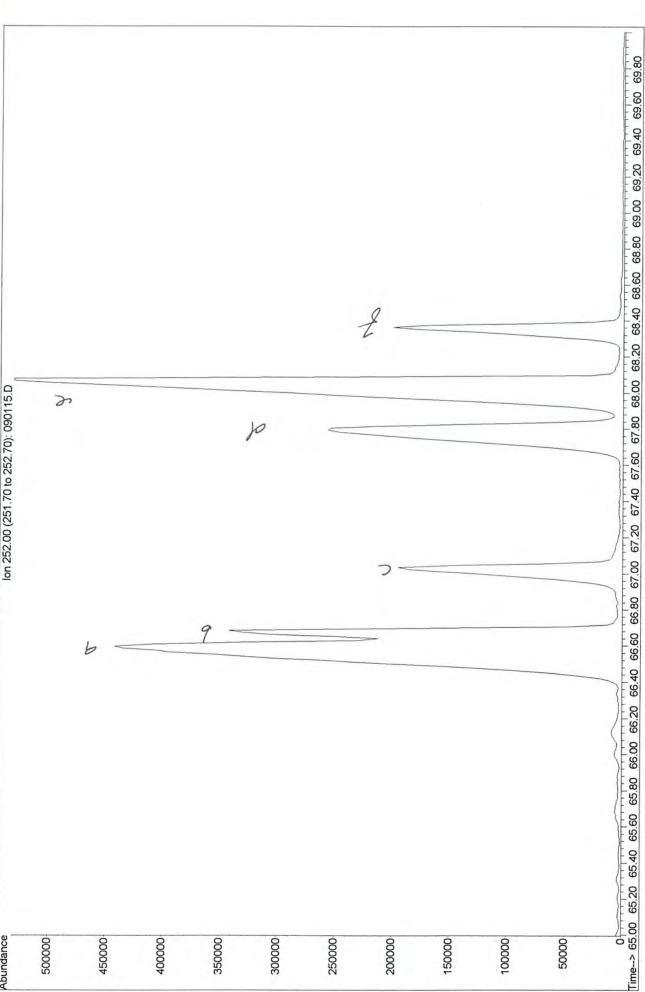


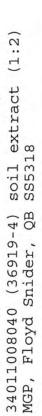


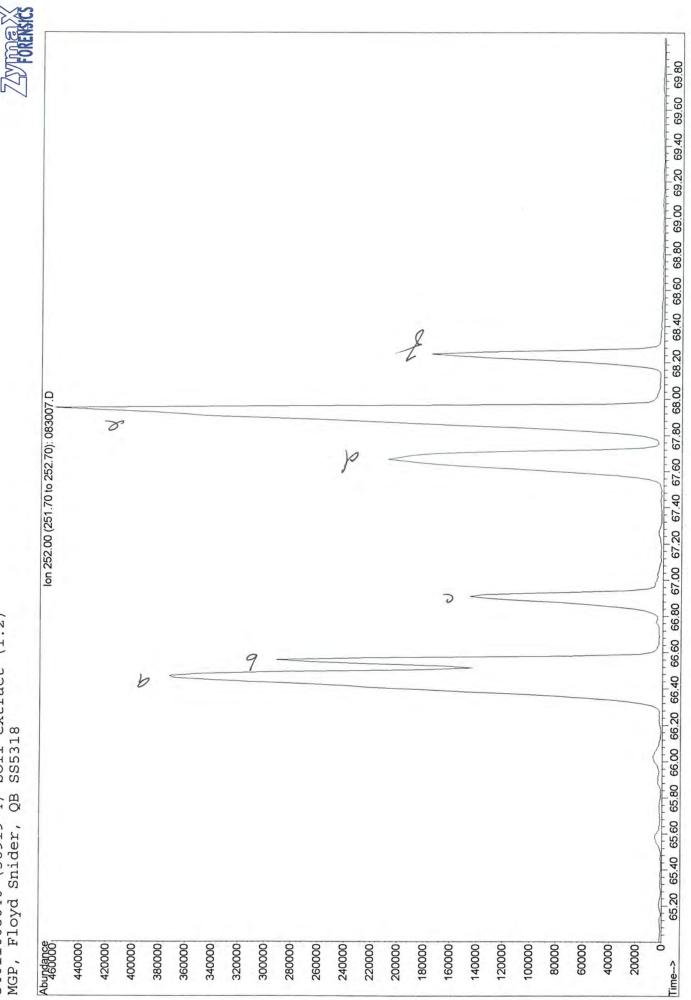
34011008032 (36918-16) soil ext (3.5:6.5) MGP, Floyd Snider, QB SS5305, Vf=4ml

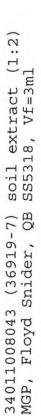


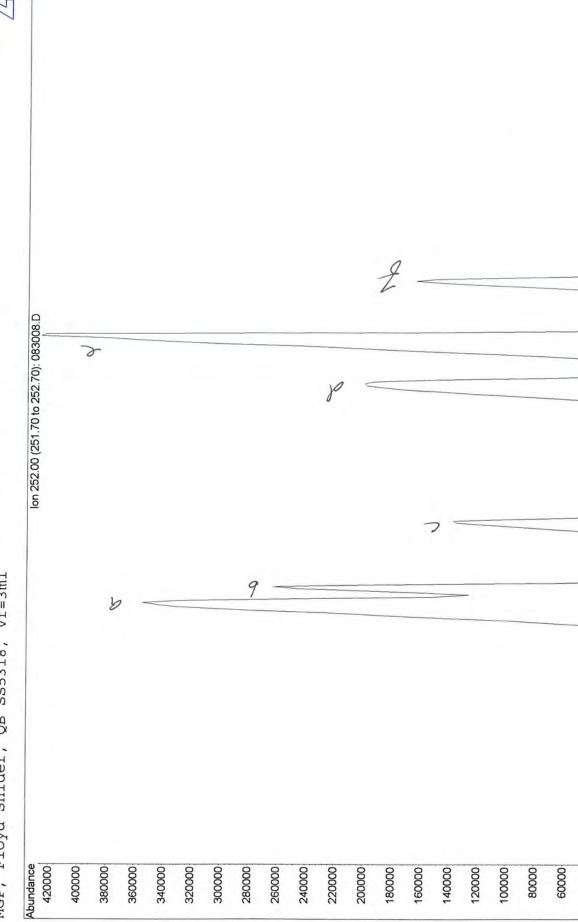










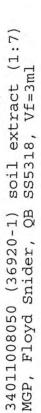


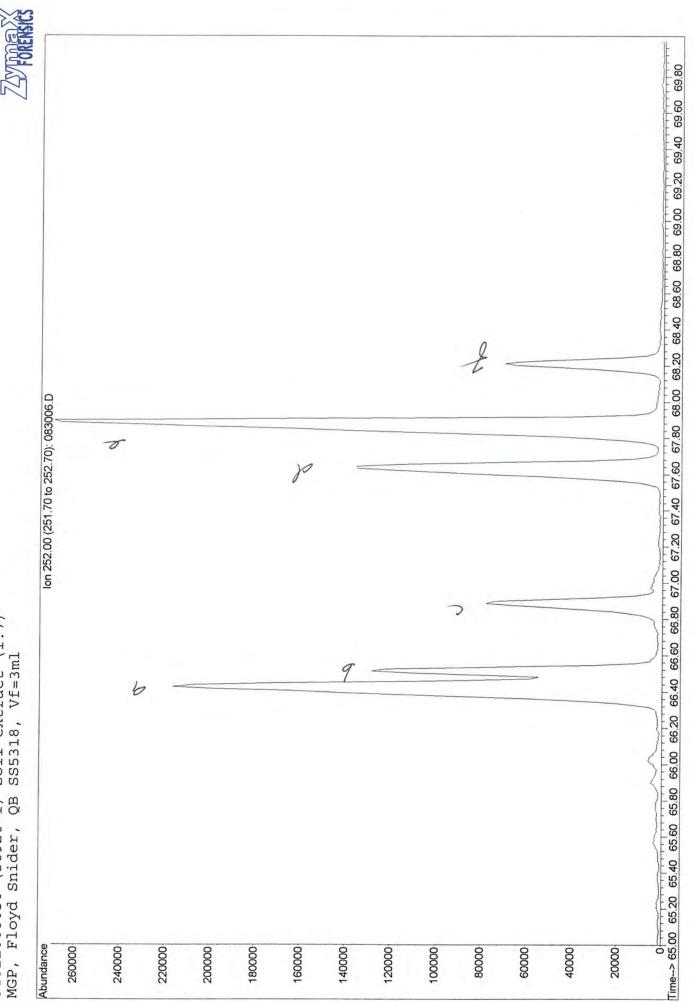
69.80

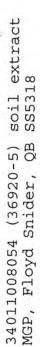
0 Time--> 65.00 65.20 65.40 65.60 65.80 66.00 66.20 66.40 66.60 67.00 67.20 67.40 67.60 67.80 68.00 68.20 68.40 68.60 68.60 69.20 69.40 69.60

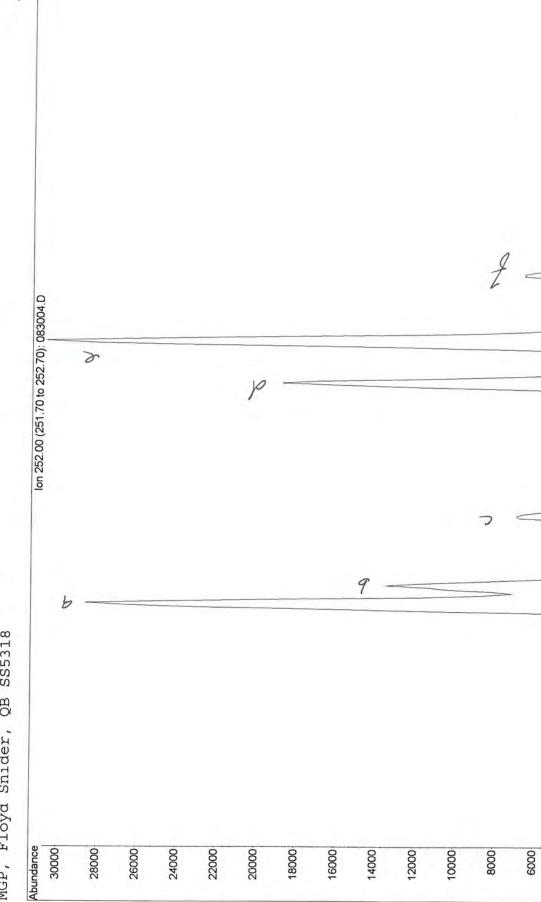
40000

20000



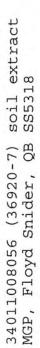


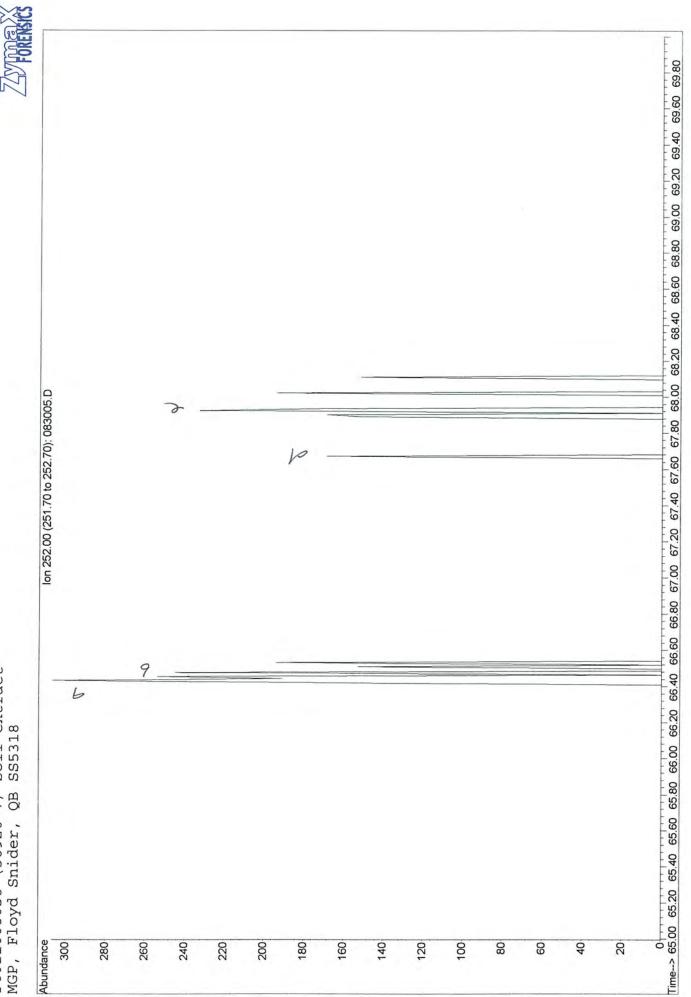


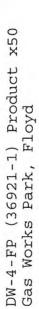


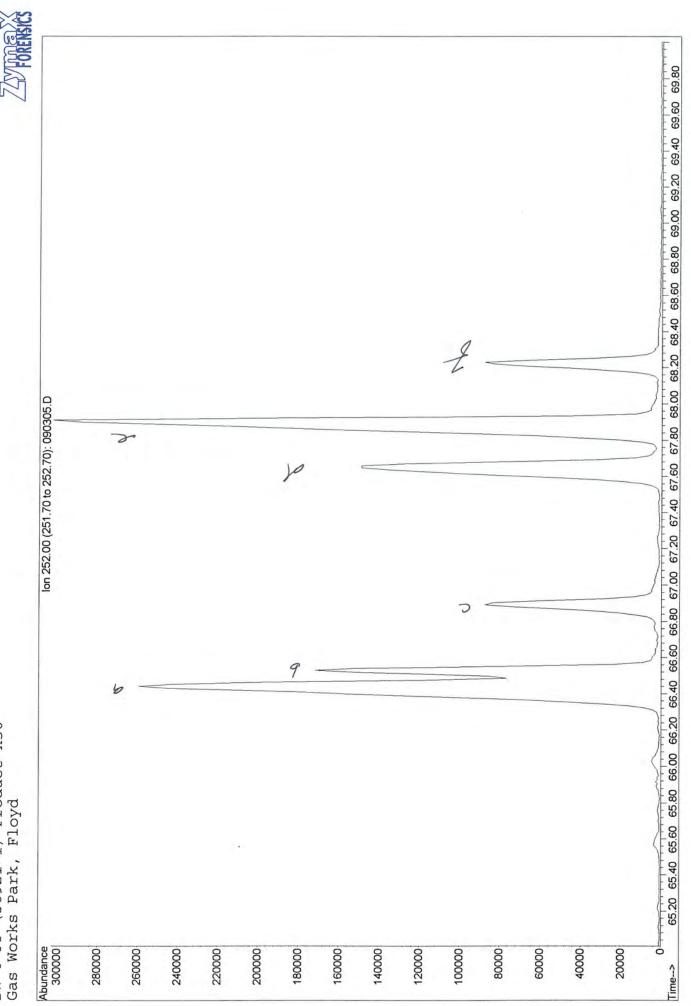
Time--> 65.00 65.20 65.40 65.60 65.80 66.00 66.20 66.40 66.60 67.00 67.20 67.40 67.60 67.80 68.00 68.20 68.40 68.60 68.80 69.00 69.20 69.40 69.60 69.60

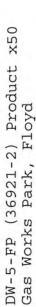




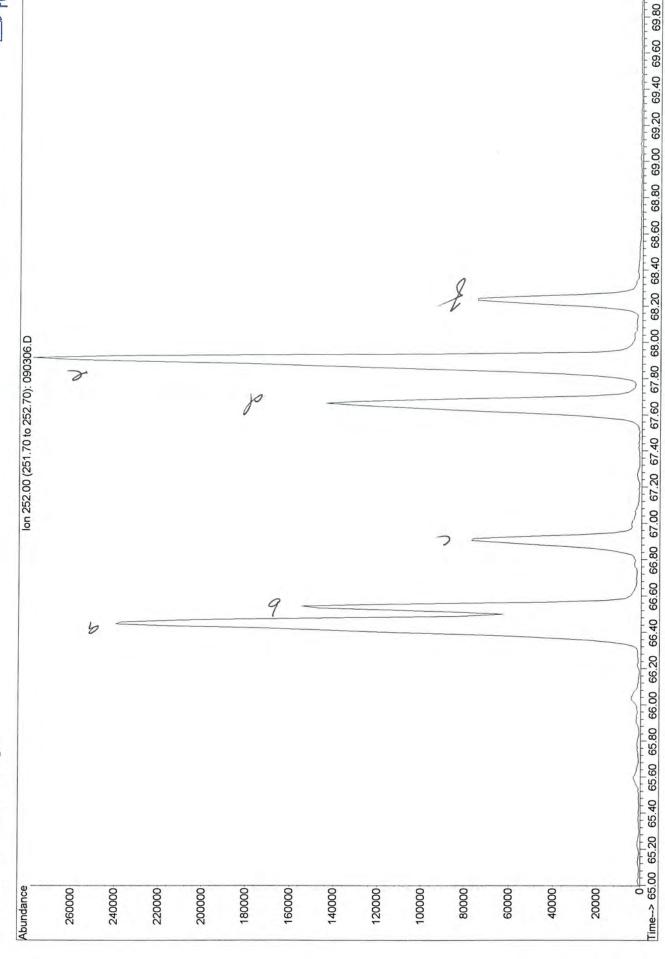


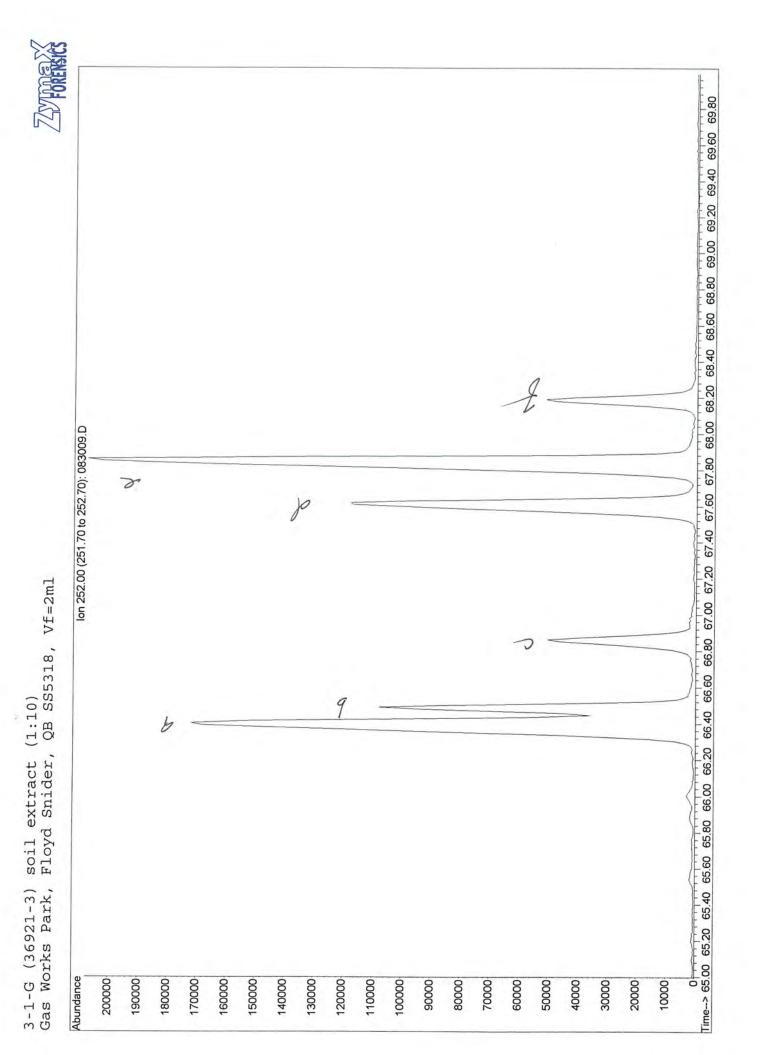


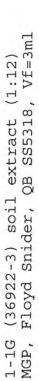


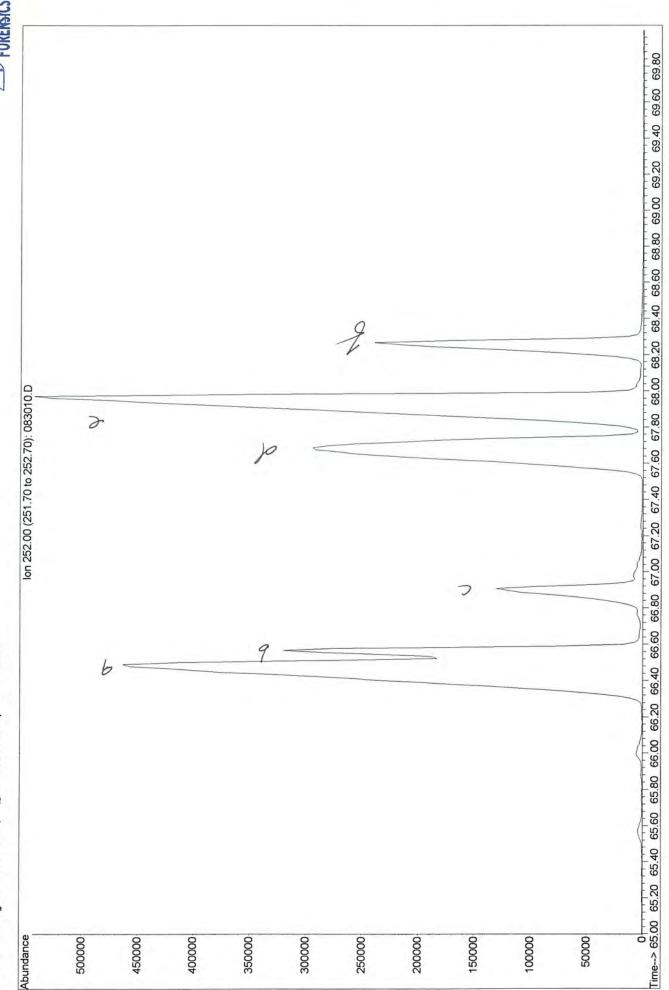


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Table

Key for Aromatic Compounds Identification in Bar Diagram

AB:	C ₃ -C ₆ Alkylbenzenes
NAPH:	C0-C4 Naphthalenes
FL:	C ₀ -C ₄ Fluorenes
BP:	C ₀ -C ₂ BP Biphenyl/Dibenzofuran
PHEN:	C_0 - C_4 Phenanthrenes
PY:	C_0 - C_4 Pyrenes/Fluoranthenes
CHR:	C ₀ -C ₄ Chrysenes
BT:	C_1 - C_5 Benzothiophenes
DBT:	C0-C4 Dibenzothiophenes
NBT:	C0-C4 Naphthobenzothiophenes
MAS:	Monoaromatic Steranes
TAS:	Triaromatic Steranes

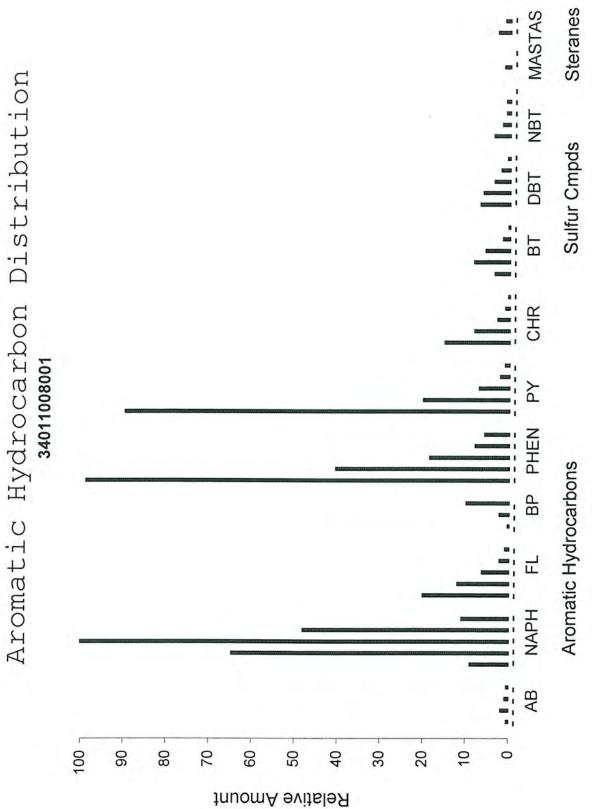
Table



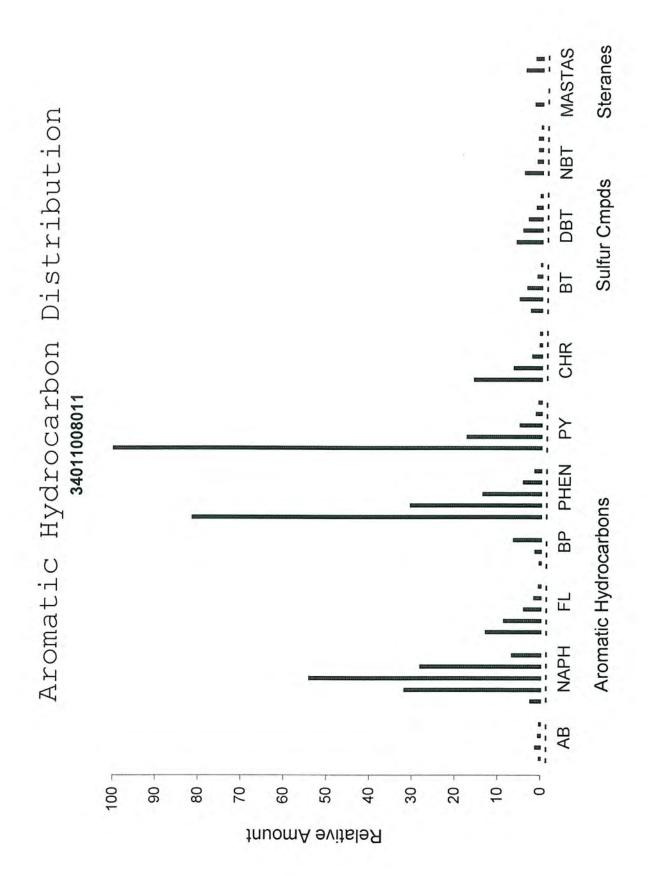
Key for Identifying Aromatic Hydrocarbons at Various m/z Units

No.	m/z	Compound	
1	120	C ₃ -alkylbenzenes	
2 3	134	C ₄ -alkylbenzenes	
3	148	C₅-alkylbenzenes	
4	162	C ₆ -alkylbenzenes	
5	128	C ₀ -naphthalene	
6	142	C ₁ -naphthalenes	
7	156	C ₂ -naphthalenes	
8	170	C ₃ -naphthalenes	
9	184	C₄-naphthalenes	
10	166	C ₀ -fluorene	
11	180	C ₁ -fluorenes	
12	194	C ₂ -fluorenes	
13	208	C ₃ -fluorenes	
14	222	C₄-fluorenes	
15	154	C ₀ -biphenyl	
16	168	C_1 -biphenyls + dibenzofuran	
17	182	C_2 -biphenyls + C_1 -dibenzofuran	
18	178	C₀-phenanthrene	
19	192	C ₁ -phenanthrenes	
20	206	C_2 -phenanthrenes	
21	220	C_3 -phenanthrenes	
22	234	C_4 -phenanthrenes	
23	202	C ₀ -pyrene/fluoranthene	
24	216	C ₁ -pyrenes/fluoranthenes	
25	230	C ₂ -pyrenes/fluoranthenes	
26	244	C ₃ -pyrenes/fluoranthenes	
27	258	C ₄ -pyrenes/fluoranthenes	
28	228	C_0 -chrysene	
29	242	C ₁ -chrysenes	
30	256	C ₂ -chrysenes	
31	270	C ₃ -chrysenes	
32	284	C₄-chrysenes	
33	148	C ₁ -benzothiophenes	
34	162	C_2 -benzothiophenes	
35	176	C ₃ -benzothiophenes	
36	190	C₄-benzothiophenes	
37	204	C₅-benzothiophenes	
28	184	C_0 -dibenzothiophene	
39	198	C ₁ -dibenzothiophenes	
40	212		
41	226	C ₂ -dibenzothiophenes	
42	240	C ₃ -dibenzothiophenes	
43		C₄-dibenzothiophenes	
+3 14	234	C ₀ -naphthobenzothiophene	
	248	C ₁ -naphthobenzothiophenes	
15	262	C ₂ -naphthobenzothiophenes	
16	276	C ₃ -naphthobenzothiophenes	
17	290	C ₄ -naphthobenzothiophenes	
18	253	Monoaromatic steranes	
19	267	Monoaromatic steranes	
50	239	Monoaromatic steranes	
51	231	Triaromatic steranes	
52	245	Triaromatic steranes	



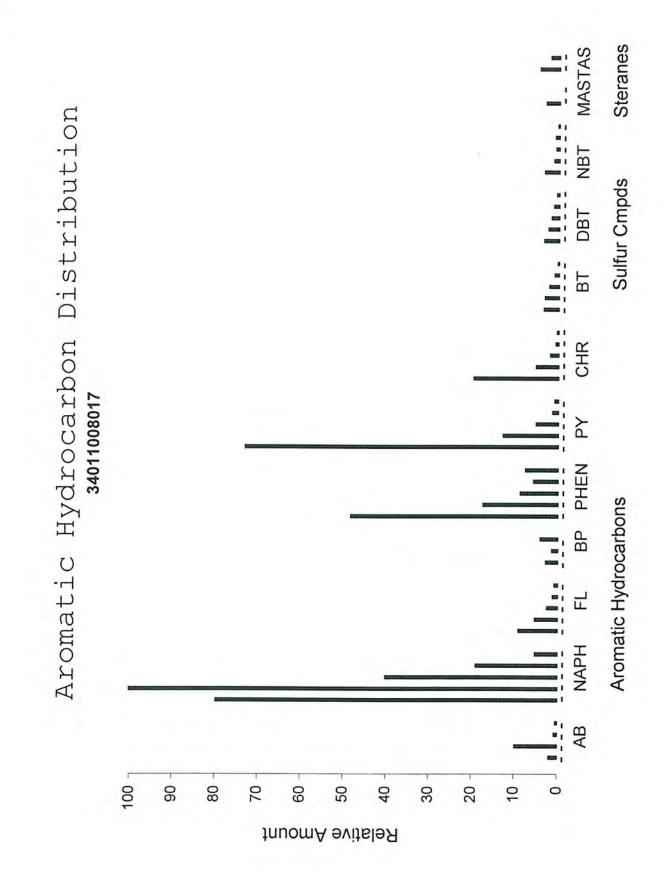




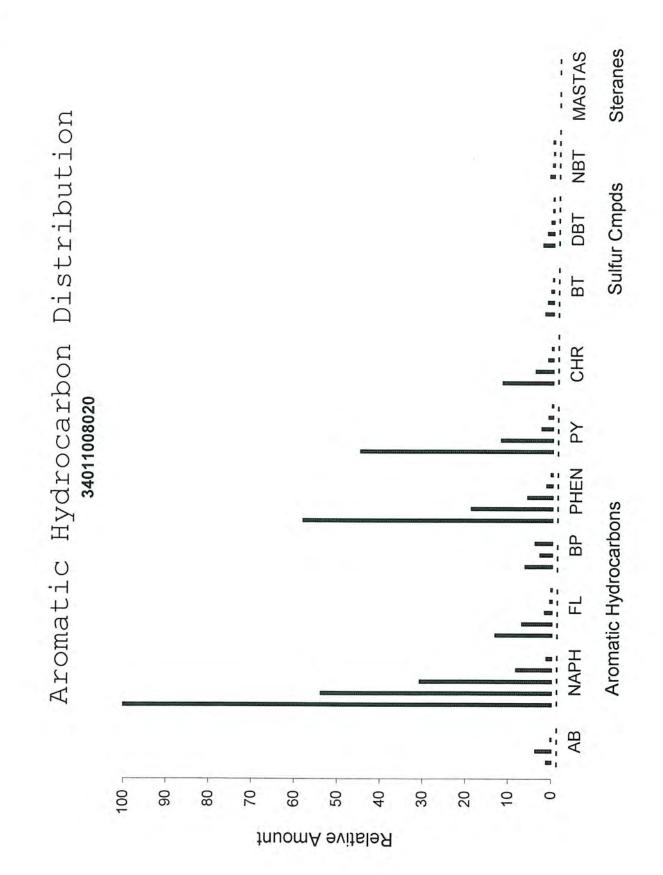


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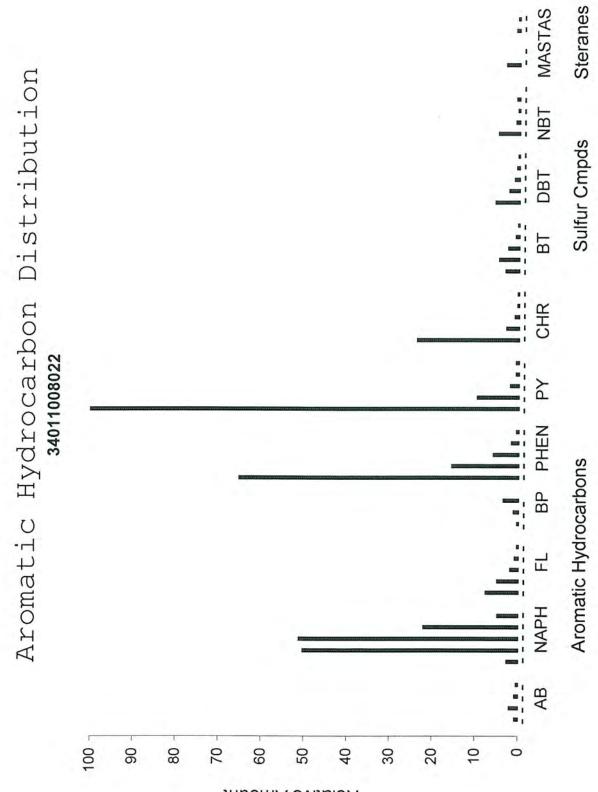






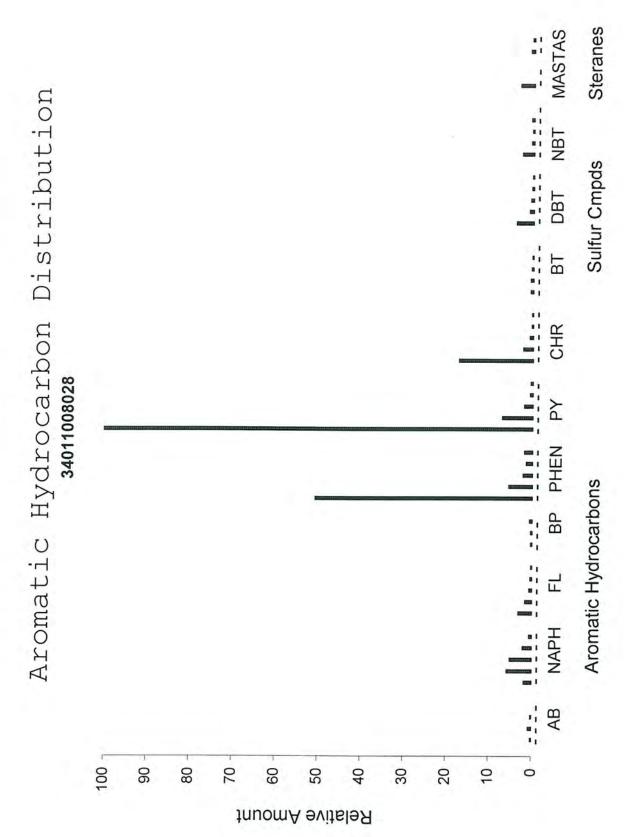




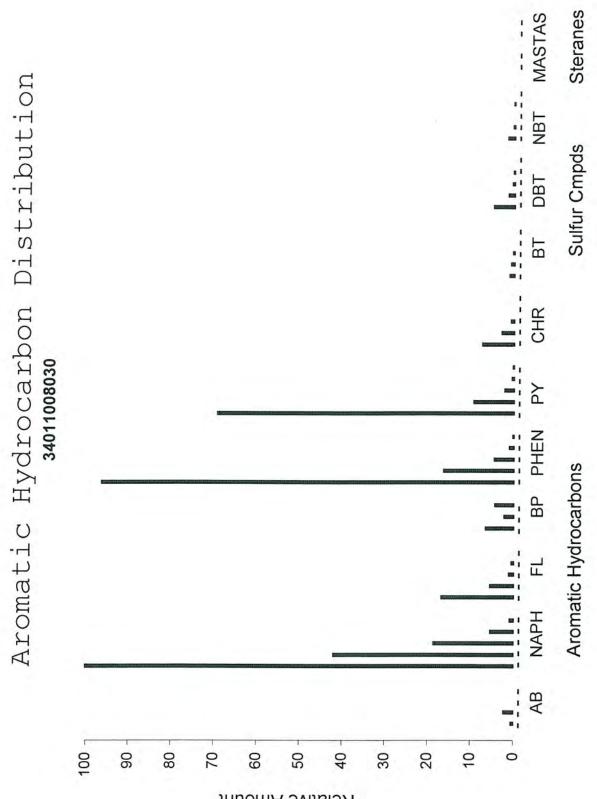


Relative Amount



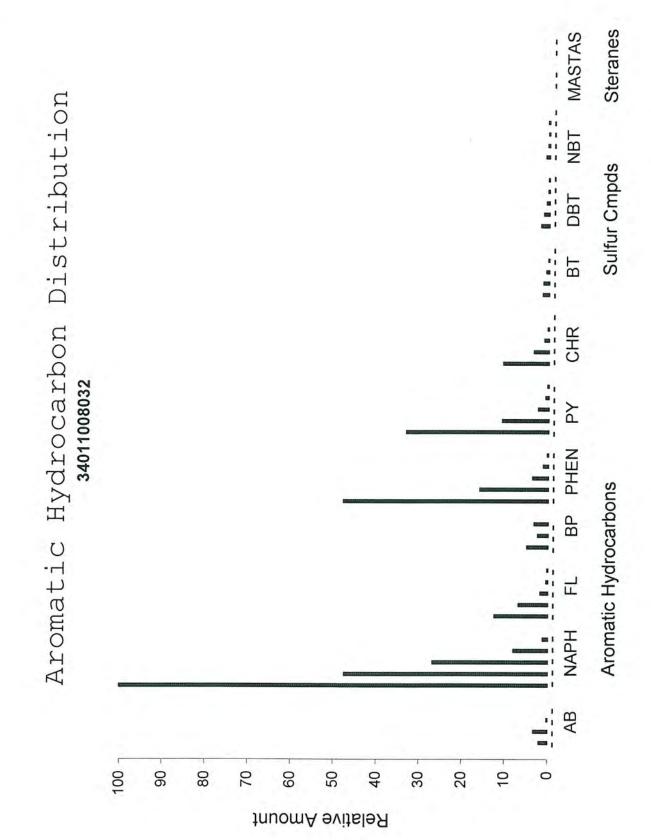




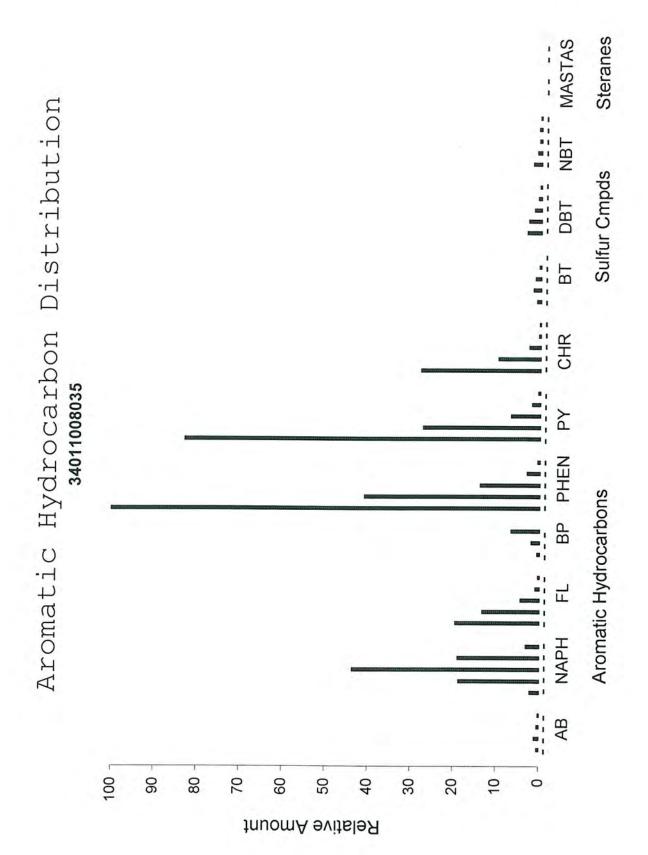


Relative Amount

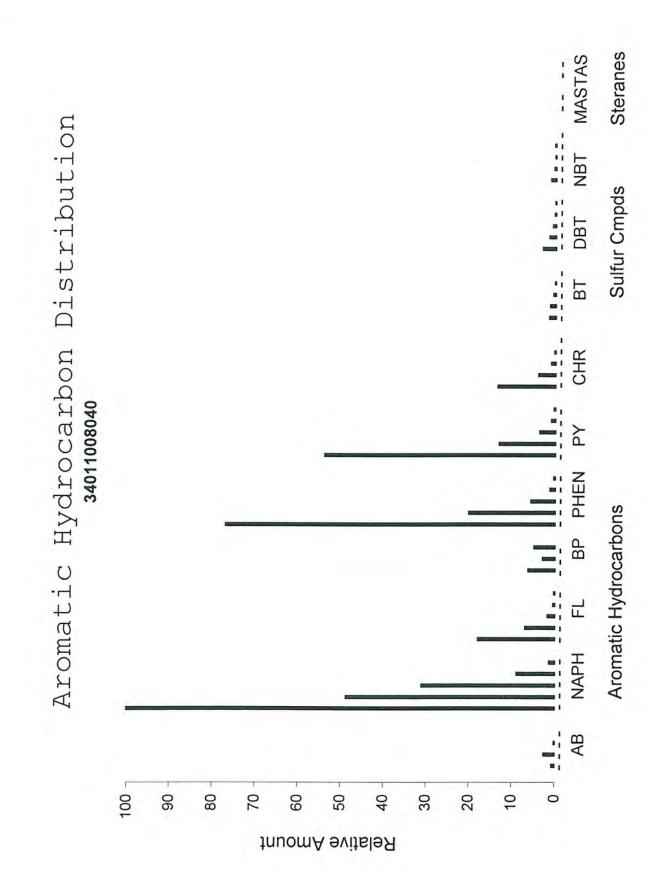




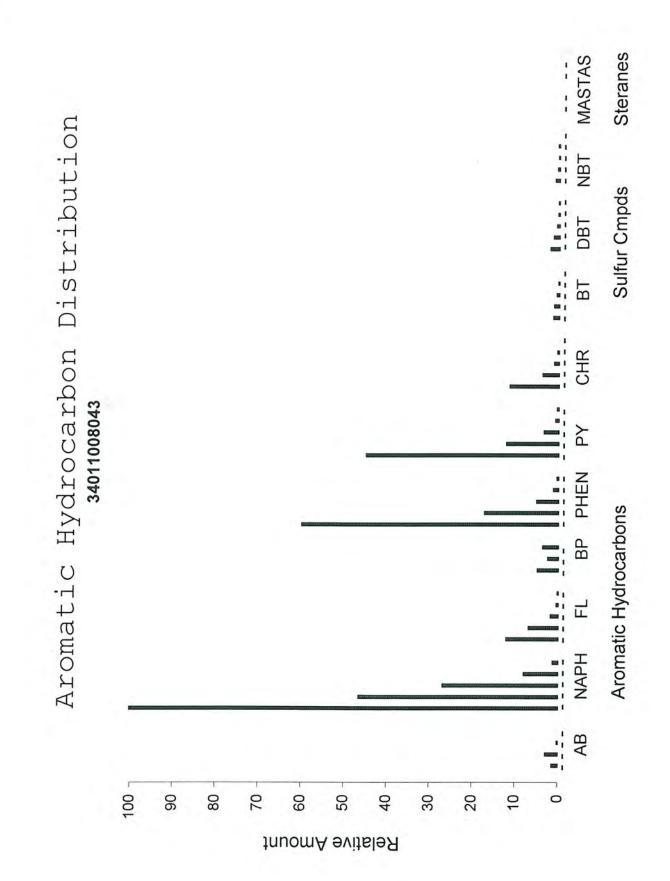






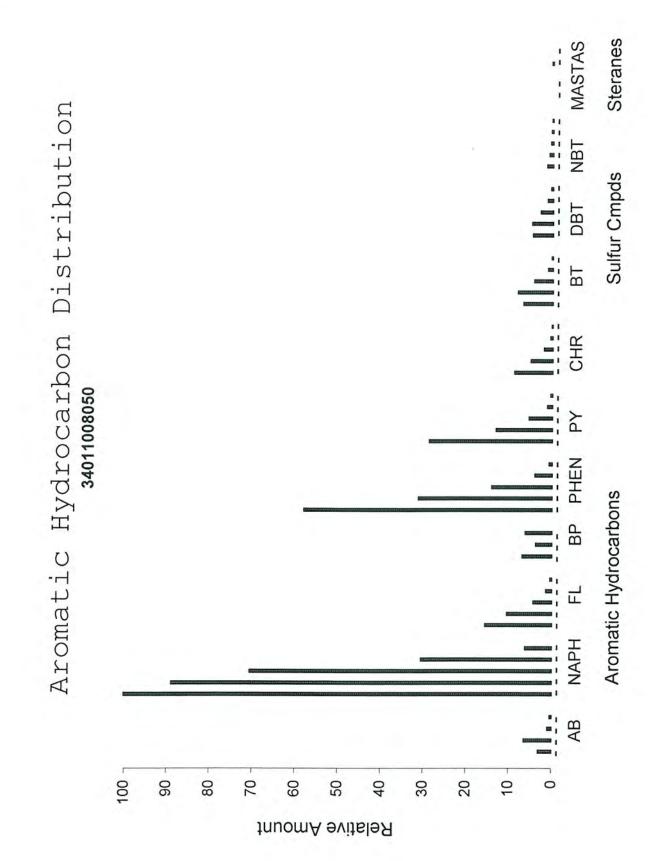




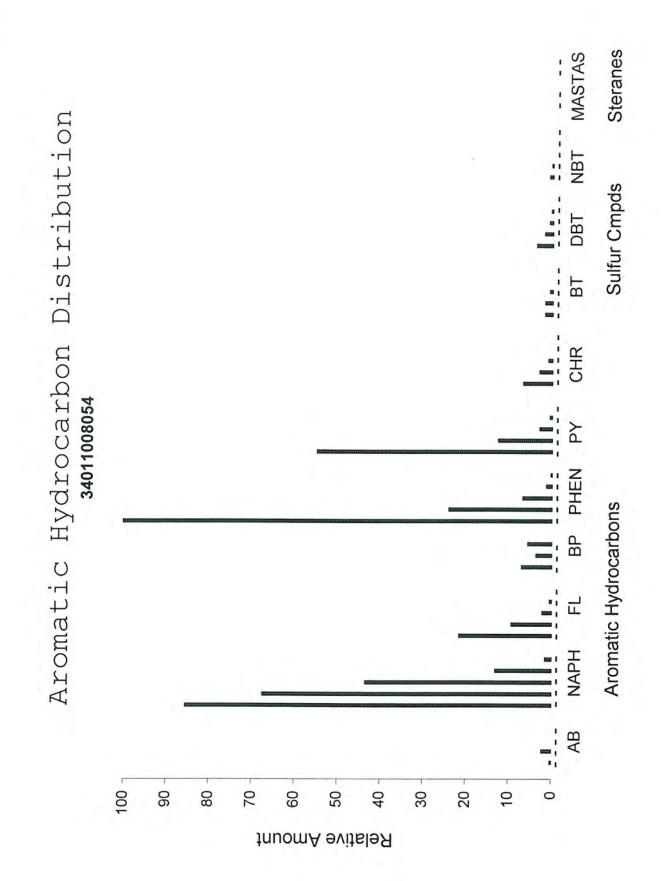


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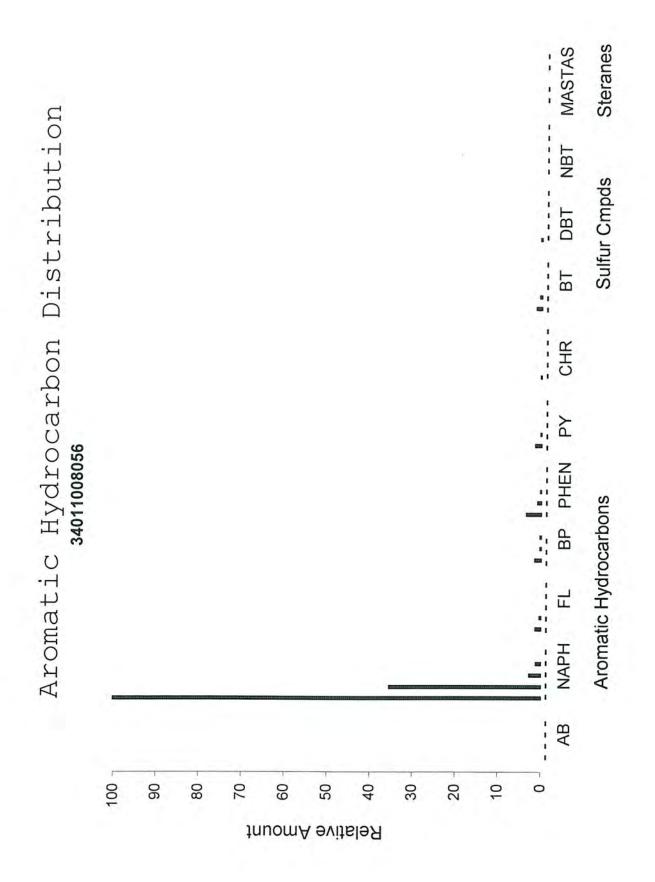




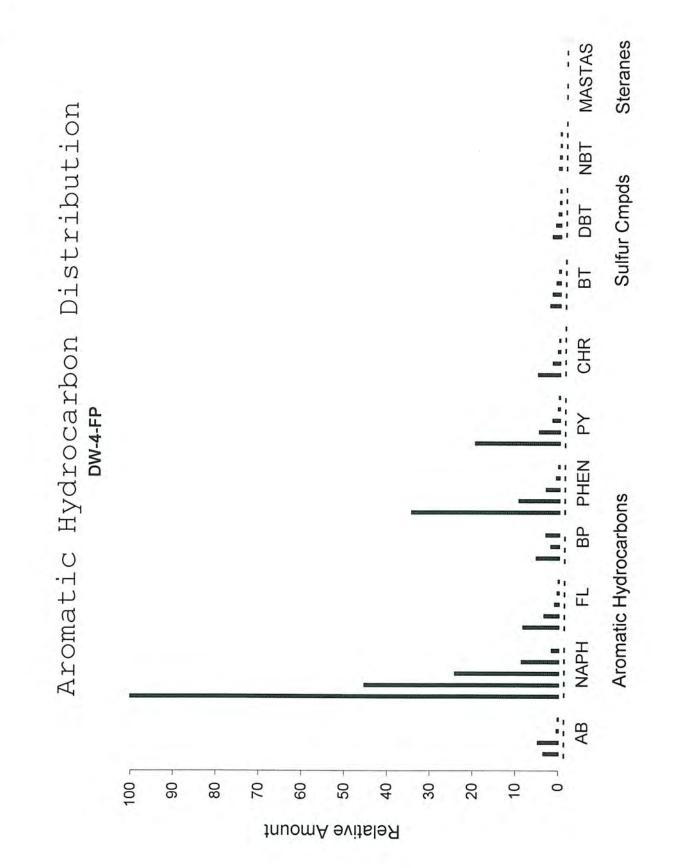




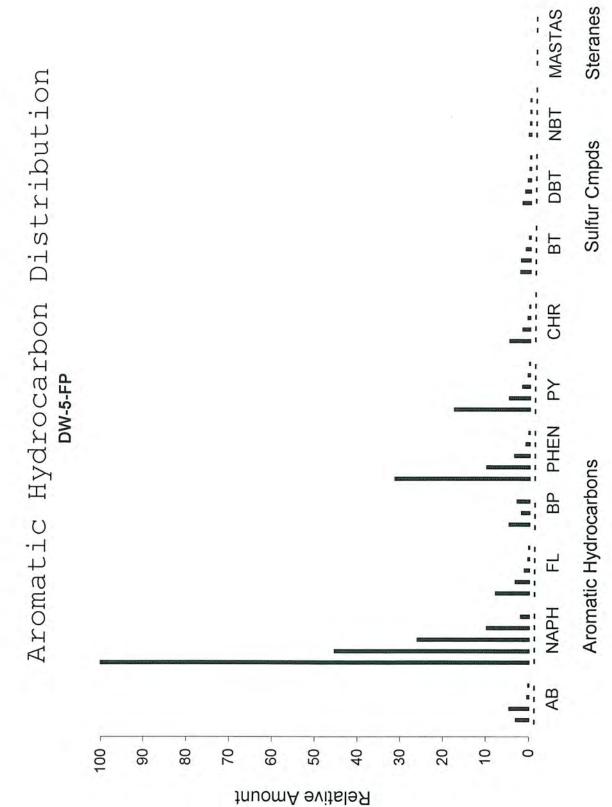






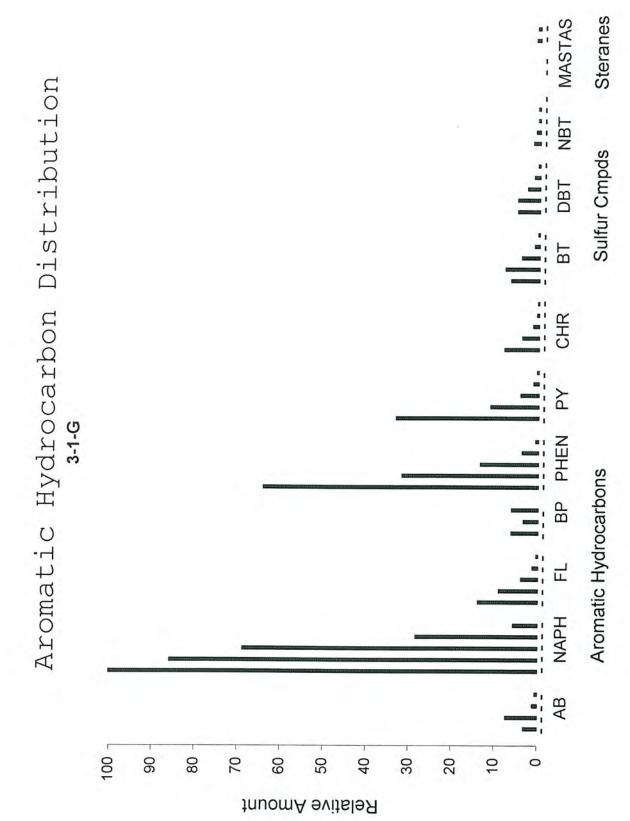




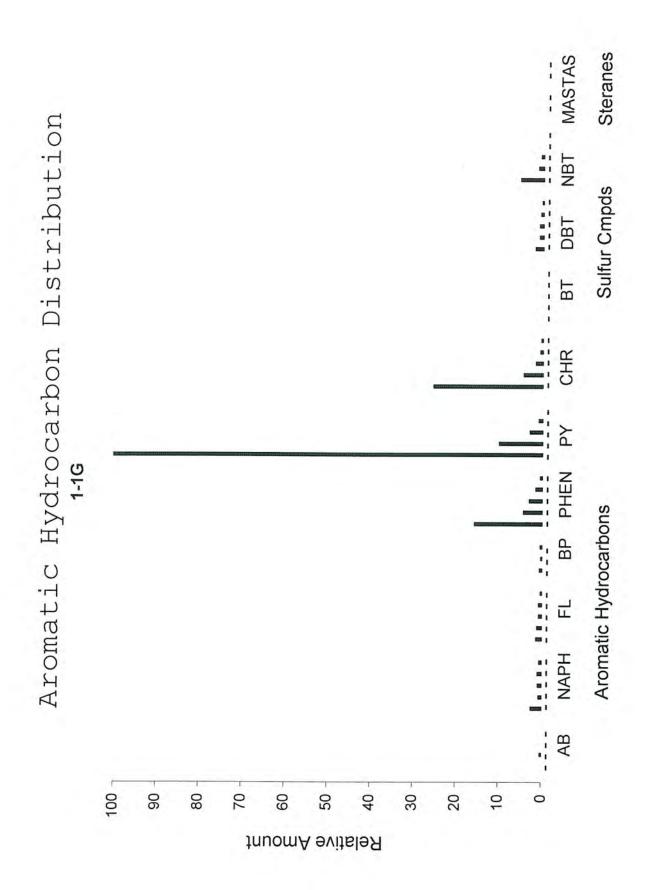


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SUB-ATTACHMENT 2D-4.2 NAPL Investigation Sediment Core Logs

Seattle Law Department Gas Works Park

Chemical Forensics Sampling Report

Appendix B NAPL Investigation 2004 Sediment Core Logs

DRAFT

CONFIDENTIAL AND PRIVILEGED: PREPARED IN PREPARATION FOR LITIGATION

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 1--1

Maximum depth of retained sediment: 7.2 ft

Mudline elevation:

8.7 ft (Corps lake datum)

	Core	Laboratory	
	collection	processing	
Date:	8/11/2004	Aug. 11, 2004	
Time:	13:40	15:00	

Percent recovery (on-deck):

74%

0 -	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
1 -	Black; mixture of water, wood chips, clay, silt, gravel and oil; More oily from 1.2 to 1.5; More wood near top, gravel near bottom; very soft, wet.	Upper recent lake		34011008028	
2	Gray; sandy gravel (GW) with f-c rounded gravel and f-c sand. Some gray, silt lumps or beds. From 1.8 to 2.1 is laminated, silty gravel (GM) with some oil penetration. No hydrocarbon				
Depth below mudline (ft.)	odor or stains below GM. Dense, moist.		Sampled GW.	34011008029	
Dept	Sediment lost during core recovery.	- - - - - - -			
6 - - - 7 -	End of Core	End of core	End of core	End of core	End of core
8 -	-				

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 1--2

Maximum depth of retained sediment: 6.9 ft

Mudline elevation: -10.8 ft (Corps lake datum)

	Core	Laboratory	
	collection	processing	
Date:	8/12/2004	Aug. 12, 2004	
Time:	10:35	14:40	

Percent recovery (on-deck): 70%

Field Log: J. LaManna

Summary Log: J. LaManna

0 -	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
	-			34011008046	
-	-	-			
	-	-			
· 1 -	Black with minor gray beds; mixture of	- Upper recent lake			
	water, wood fibers and fragments, clay and oil. Very soft, wet.	deposits			
-	-	-			
-	-	Ì			
2 -					
-	-				
-	-	-			
-	-	ļ			
3 -					
(ft.)	Gray interbedded sandy gravey (GW),	-			
line	sand (SP) and sandy silt (SM). GW 6.7- 7.6, ML+SM 7.7-8.3, SP 8.3-9.0, SM	Stratified Drift			
pnu	9.0-9.1. Dense, wet. No oil, sheen or stains.	-			
N 4 -					
belo	+				
Depth below mudline (ft.)	-	-	Sampled SP.	34011008045	
	-	-			
5 -					
-	-	-			
	-	-			
6 -	- Sediment lost during core recovery	I			
	-	+			
	-	+			
	† I	1			
7 -	End of Core	End of core	End of core	End of core	End of core
-	-	ł			
-		1			
-	-	+			
8 -		<u> </u>			

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T3

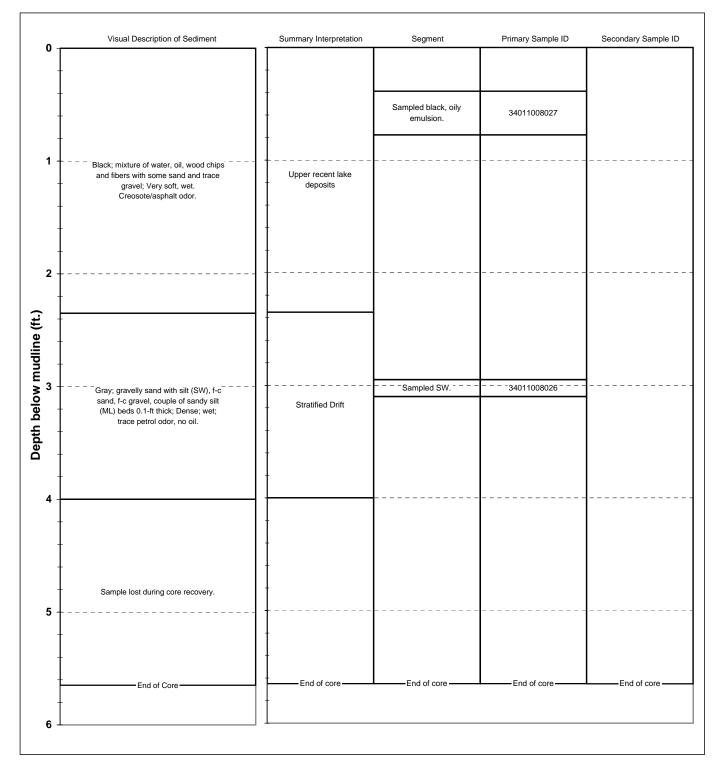
Station: 1--3

Maximum depth of retained sediment: 5.7 ft 74%

Mudline elevation: 20.9 ft (Corps lake datum)

Percent recovery (on-deck):	7

	Core	Laboratory
	collection	processing
Date:	8/11/2004	Aug. 11, 2004
Time:	12:35	14:45



Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 2-1 Rep 2

Maximum depth of retained sediment: 6.1 ft

-12.7 ft (Corps lake datum) Mudline elevation:

	Core	Laboratory
	collection	processing
Date:	8/11/2004	Aug. 11, 2004
Time:	17:00	18:00

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58%

Field Log:	J. LaManna
Summary Log:	J. LaManna

0 -	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
• • •	Dark olive; mixture of wood fragments (some lumber scraps), silt, clay and some oil. Very soft (almost a liquid), wet. Smells like creosote or asphalt.	Upper recent lake deposits		34011008036	
1 - - -	Dark olive; silty clay (CL) with black oil stains from 0.9 to 1.1 and black lamellae throughout. Very soft, wet.	Lower recent lake		34011008035	
2					
Depth below mudline (ft.)					
Depth bel	Gray; sandy gravel (GW) with f-c sand and f-c rounded gravel, c gravel and cobbles at 5.2; Dense, wet; no oil or stains.	- Stratified Drfit - -		34011008034	
5 -		-			
6 -	End of Core	End of core	 End of core	End of core	End of core
7 -	-				

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 2-2 A

Maximum depth of retained sediment: 9.7 ft

Mudline elevation: -17.0 ft (Corps lake datum)

	Core	Laboratory	
	collection	processing	
Date:	8/12/2004	Aug. 12, 2004	
Time:	9:00	12:00	

Percent recovery (on-deck): 72%

0 -	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
2 -	Black with minor dark gray; mixture of water, clay, wood fragments and tarry oil; Very soft, wet.	- Upper recent lake deposits			
-	-	-	— Sampled gray sitty clay — — with black tarry oil. —	34011008043	
4 - -	+	+ 			
mudline (ft.)	-	-	Sampled SW with bright sheen and black oil.	34011008042	
Depth below mudline (ft.)	 Gray; interbedded sandy gravel (GW) and gravelly sand (SW) with SP from 5.0-6.4, brown wood fibers abundant in upper part of interval; Dense, wet. Brightly colored sheen with black oil 3.6-4.3 and 7.8-8.6; 	Stratified Drift	 Sampled SP with no visible oil. 	34011008041	
- 10 - 8		- - 	Sampled GW with bright sheen and black oil.	34011004040	34011008044
-		- - -			
10 -	End of Core	End of core	End of core	End of core	End of core ———
-		-			
12 -					

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T3

Station: 2--2

Maximum depth of retained sediment: 9.7 ft

Mudline elevation: -14.3 ft (Corps lake datum)

	Core	Laboratory
	collection	processing
Date:	8/12/2004	Aug. 12, 2004
Time:	9:33	11:00

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Percent recovery (on-deck): 82%

0 -	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
	Black; mixture of water, clay, wood and oil; Very saft, wet; black oil blebs and slight sheen; asphalt odor.	- Upper recent lake - deposits		34011008039	
2 -					
-	Blotched or mottled, black and dark gray; organic clay (OL); some wood fragments near core tube may be carry down; very soft, wet; black may be oil.	Lower recent lake		34011008038	
4 -					
Depth below mudline (ft.) 9	Gray; interbedded sandy gravel (GW) and gravelly sand (SW) with a sitly clay (CL) from 5.6-5.9 and gravelly sitt (ML) below 7.9; Dense, wet; faint hydrocarbon odor; oil stains and sheen from 4.0-4.5.	- - - - - - Stratified Drift	— Sampled sand with few — — fines below silty clay. —	34011008037	
8 -					
10 -	End of Core	End of core	End of core	End of core	End of core ———
12 -	-				

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 3-3 Rep 3

Maximum depth of retained sediment: 14.0 ft 72%

Mudline elevation: -18.6 ft (Corps lake datum)

	Core	Laboratory		
	collection	processing		
Date:	8/11/2004	Aug. 11, 2004.		
Time:	15:07	16:30		

•	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
0	Black; mixture of water, wood chips, organic clay, some rounded gravel and black oil; Very soft, wet. Creosote or asphalt odor, very oily.	Upper recent lake deposits		34011008033	
2		* 			
4 +	Dark olive gray; organic clay (OL), with veins of black oil from 1.1-5.8 and 6.9- 7.5; Very soft, wet. Black oil stains from 1.1 to 1.3.	Lower recent lake deposits			
Idline (ft.)			Sampled oily vein.	34011008032	
Depth below mudline (ft.) * + + + + + + + + + +	Dark olive gray viscous liquid that appears to be mixture of organic clay and black tarry oil; very soft; wet.	- - - -	Sampled oily GW	34011008031	
10 +	Gray; interbedded sandy gravel (GW), gravelly sand (SW) and below 11.3 minor silty sand (SM); Dense, wet. Black tarry oil in sandy beds 8.8-10.0 and 10.7-10.9. Smells like creosote or asphalt.	Stratified Drift	Sampled oily GW.	34011008030	
12 +	Sediment lost during core recovery	 - - -			
14	End of Core	End of core	End of core	End of core	End of core
16		+			

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 3-5 Rep 2

Maximum depth of retained sediment: 5.7 ft

Mudline elevation: 20.9 ft (Corps lake datum)

Percent recovery (on-deck): 29%

 Core
 Laboratory

 collection
 processing

 Date:
 8/13/2004
 Aug. 13, 2004

 Time:
 17:30
 13:45

Field Log: J. LaManna

Summary Log: J. LaManna

•	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
0	Black grading down to gray; mixture of water, clay, wood and black tarry oil;	Upper recent lake deposits	Sampled gelatinous tarry material.	34011008058	
	Very soft, wet. Asphalt odor.				
	+	+			
	1				
	†	1			
1 -	+	+			
	1	+			
	Dark olive gray; organic clay (OL) with a piece of wood; Very soft, wet. Black	Lower recent lake			
	oil-stain from 1.7 to 1.8	- deposits			
	+	+			
	-	+	Sampled black, oil-		
2			stained OL.	34011008057	
2	Τ				
	+	+			
ft.)	+	-			
) el		-			
alir			Sampled oily GW.	34011008056	
nu	†				
Depth below mudline (ft.) $_{\omega}^{\circ}$	+	+			
elo	+	-			
d d	Dark gray; gravelly sand (SW) grading				
<u>s</u> ptl	down to sandy gravel (GW); Dense, wet. Oil stains at 2.7.				
ð		Ť			
	+	+			
4					
-		Stratified Drift			
	†	1			
		-			
	-	+			
	†				
5	Sediment lost during core recovery.	•			
	ļ				
	† I				
	End of Core	End of core	End of core	End of core	End of core
			3, 5010	2. 0010	
6					
σ					

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 3--1

Maximum depth of retained sediment: 4.4 ft

Mudline elevation: -3.5 ft (Corps lake datum)

	Core	Laboratory
	collection	processing
Date:	8/11/2004	Aug. 11, 2004
Time:	10:42	14:00

Percent recovery (on-deck): 82%

0 -	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
-	- - - -				
1 - - -					
1 - - -		+ - - - -			
2 -	Black; mixture of water, wood chips, silt, clay, sand, oil and trace gravel; most wood chips in top 0.3-ft; most silt and sand near bottom; Very soft (gelatinous to liquid in places), wet; smells like creosote or asphalt.	Upper recent deposits			
Depth below mudline (ft.)		+ - - - -			
pth below		+ - - - -			
۳ ۵ - - -		+ - - -	− − − Sampled very oily − − − interval.	34011008025	
4 - - -					
- 4 - -	Sediment lost during core recovery.	+ 			
5 -	End of Core	End of core	End of core	End of core	End of core
5 -		<u> </u>			

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 3--2

Maximum depth of retained sediment: 9.3 ft 69%

20.9 ft (Corps lake datum) Mudline elevation:

•				
Percent	recovery	(on-deck)):	6

	Core	Laboratory	
	collection	processing	
Date:	8/11/2004	Aug. 11, 2004	
Time:	8:30	11:00	

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Field Log: J. LaManna

Summary Log: J. LaManna

	0 -	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
		-	-		34011008017	
	- - 1 - - -	Black grading down to dark greenish gray; sandy silt (ML) with wood chips and one shell; Very soft, wet. Black oil and sheen.	Upper recent lake			
	2 -					
	3 -	Dark greenish gray; sandy gravel with silt (GW) or gravelly sand with silt (SW) with wood chips and root fibers; Very soft, wet. Black oil blebs in most of interval, oil coats root fibers; strong asphalt odor.	Upper recent lake - deposits; possibly Gas Works Fill		34011008018	
ft.)	4 -		+			
line (-	-	+	Sampled oily SW.	34011008020	
Depth below mudline (ft.)	5 -	- - 	- - - -			
Jepth be	6 -	Dark gray; gravelly sand (SW) grading down to sand with gravel (SP); Dense, moist. Asphalt odor, some oil stuck on gravel.	- - Stratified Drift			
-	-	-	-			
	7					
	-	-			34011008019	
	8 -	Sediment lost during core recovery; core tube bent and scratched.				
	9 -	End of Core	End of core	End of core	End of core	End of core
	10		-			

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 3--3

Maximum depth of retained sediment: 7.2 ft

-18.5 ft (Corps lake datum) Mudline elevation:

	Core	Laboratory
	collection	processing
Date:	8/11/2004	Aug. 11, 2004
Time:	9:40	13:00

Percent recovery (on-deck): 71%

0 -	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
0 -		-	Sampled black, gelatinous, oily material.	34011008022	
+ + + 1 -	Gray (top 0.1-ft) and black mixture of clay, water, silt, wood fibers (possibly bark) and oil; Very soft, wet. Heavy oil 0.0-0.2 and 0.9-1.2; black oil appears to penetrate next lower interval	Upper recent lake deposits			
-	· · · · · · · · · · · · · · · · · · ·	-	Sampled black oily mud.	34011008024	
2		-			
2 -					
-	· · · · · · · · · · · · · · · · · · ·	-			
(ff.)	· · · · · · · · · · · · · · · · · · ·				
v mudline	Dark olive gray; organic clay (OL) with trace fir needles; Very soft, wet.	Lower recent lake			
Depth below mudline (ft.)	abundant at top of interval; oil content highest at top and bottom of interval. Did not observe veins from 3.8-4.5.	deposits			
5 -	· · · · · · · · · · · · · · · · · · ·				
-		-	Sampled black vein.	34011008023	
6 -	· · · · · · · · · · · · · · · · · · ·	 			
7	Black oily wood chunks; very little sediment. May be repenetration of surface material. However, boat crew	Provenance not	Sampled black, viscous oil.	34011008021	
'	reports refusal. End of Core	End of core	End of core	End of core	End of core ———
		+			
8		<u> </u>			

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 3--4

Maximum depth of retained sediment: 10.5 ft

Mudline elevation: -13.8 ft (Corps lake datum)

	Core	Laboratory
	collection	processing
Date:	8/12/2004	Aug. 13, 2004
Time:	15:32	12:30

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Percent recovery (on-deck):

59%

⁰ ⊤	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
- - - 2 - -	Black grading down to dark gray; clay (CL) with water, wood fragments, oil and with black, angular gravel-sized grain that, crushed between the 	Upper recent lake deposits	Sampled black oil.	34011008055	
4 +			Sampled oily GW with bright sheen.	34011008054	
6 + + + +	Dark gray sandy gravel (GW), f-c gravel, top of interval more sandy; Dense, wet. Black oil with bright sheen at 4.3; no oil in lower portion.	- Stratified Drift			
8 -	Gray sandy gravel (GW). About 50% recovery and slumped in core tube.	-			
10 -	Sediment lost during core recovery.	End of core	End of core	End of core	End of core
12		-			

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 4-1 A

Maximum depth of retained sediment: 7.4 ft

Mudline elevation: -3.7 ft (Corps lake datum)

Percent recovery (on-deck): 84%

	Core	Laboratory		
	collection	processing		
Date:	8/10/2004	Aug. 10, 2004		
Time:	15:40	18:30		

Field Log: J. LaManna

Summary Log: J. LaManna

0 -	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
1	Black; silt with fine sand (ML) and abundant wood fragments; Very soft, wet. Oily with asphalt odor.		Sampled black, gelatinous oily material.	34011008013	
2	Dark olive interbedded silt with fine sand (ML) and sandy silt (SM) with wood chips and fibers and a glass bottle at bottom of interval; Very soft, moist-wet. Oil and sheen 1.7-2.0.	Upper recent lake deposits	Sampled oil stains.	34011008014	
Depth below mudline (ft.)	Dark gray fine-medium sand (SP) with rounded fine gravel beds (SW) 0.1-ft	Stratified Drift	Sampled oily SW.	34011008015	
□ 5 - 6 -	thick at 3.4 and 4.6; Dense, moist. No odor, sheen or stains.	* *			
7 -	Sediment lost during core recovery.	End of core	End of core	End of core	End of core

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 4-2 Rep 1

Maximum depth of retained sediment: 1.8 ft

Mudline elevation: -15.4 ft (Corps lake datum)

	Core	Laboratory
	collection	processing
Date:	8/10/2004	Aug. 10, 2004
Time:	16:48	19:00

Percent recovery (on-deck): 32%

	Visual Description of Sedi	ment Summary Interpretatio	n Segment	Primary Sample ID	Secondary Sample ID
	0 Black; silty fine (SM) or sandy (ML), with clay abundant wood verysoft, wet. 0	silt and Upper recent lake chips; deposits Oily	Sampled black gelatinous oily material.	34011008016	
(ft.)	1				
Depth below mudline (ft.)	1 Sediment lost d				
Dept	1				
	2End of Core	End of core	End of core	End of core	End of core
	2				

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 4-3 Rep 1

Maximum depth of retained sediment: 14.8 ft

Mudline elevation: -19.6 ft (Corps lake datum)

Percent recovery (on-deck): 86%		Percent recovery	(on-deck):	86%	
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	Core	Laboratory
	collection	processing
Date:	8/10/2004	Aug. 10, 2004
Time:	14:20	17:00

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0 -	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
2 -	Black grading down to dark gray; organic clay with silt, wood chips and black oil blebs; Very soft, wet. Asphalt odor.	Upper recent lake deposits		34011008011	
4	-	- - - - -			
6	- - -	+ - - - -			
Depth below mudline (ft.)	 Very dark brown; organic clay (OL);Very soft, wet. No oil or stains	- - - Lower recent lake deposits			
Depth be 10	- - - 	+ - - - -			
12 -	- - - 	+ + 			
14 -	- - - 	End of core	End of core	End of core	
16	-	-			

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

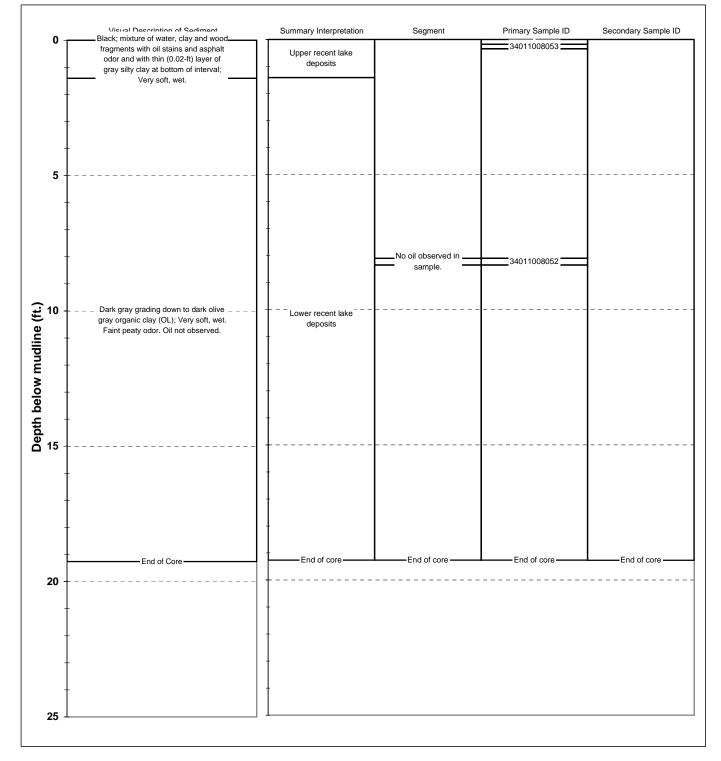
Station: 4--4

Maximum depth of retained sediment: 19.3 ft

Mudline elevation: -19.6 ft (Corps lake datum)

	Core	Laboratory
	collection	processing
Date:	8/13/2004	Aug. 13, 2004
Time:	14:18	11:45

Percent recovery (on-deck): 69%



Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

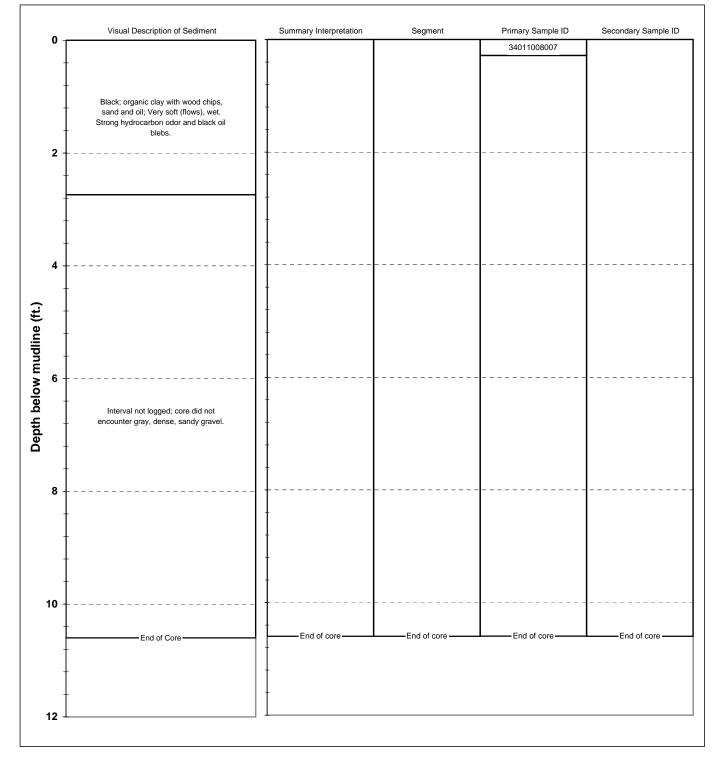
Station: 5-1

Maximum depth of retained sediment: 10.6 ft

Mudline elevation: -18.6 ft (Corps lake datum)

	Core collection	Laboratory processing
Date:	8/10/2004	Aug. 10, 2004
Time:	9:00	15:00

Percent recovery (on-deck): 79%



Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 5-1 Rep 2

Maximum depth of retained sediment: 16.0 ft

Mudline elevation: -18.1 ft (Corps lake datum)

Percent	recoverv	(on-deck):	68%	
		(0	00/0	

	Core	Laboratory		
	collection	processing		
Date:	8/10/2004	Aug. 10, 2004		
Time:	9:38	13:30		

0	Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
	Black; mixture of water, organic clay, silt and wood chips; Very soft, wet. Oil sheen and asphalt or creosote odor.	-		34011008001	
2 +	Dark gray silt (ML) with wood fragments and fibers and trace angular sand; Very soft, wet. Trace sheen	Upper recent lake deposits		34000118002	
+	Very dark brownish gray grading down to dark brown; organic clay (OL) with gray lamellae and with brown peaty lumps; Very soft, wet. No odor or	-		34011008003	
4	sheen.			34011008004	
6 +		- - 			
e (ft.)	Olive brown; organic clay (OL) with trace roots; Very soft, wet. No sheen, slight reducing odor.	- - -			
Depth below mudline (ft.) 01 8 11 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1		Lower recent lake deposits -			
bth belov					
+	Very dark brown organic clay (OL) with wood fragments and beds of gray fine	-		34011008005	
12 + -	 to medium sand 0.05 to 0.1-ft thick; Very soft and very loose; wet. No odor or sheen. 				
14	Gray; well graded, sandy gravel (GW) with f-d sand and f-c gravel, with lumps of sandy silt (ML); loose, moist. No hysrocarbon odor or sheen.	- Stratified Drift		<u></u> 34011008006 <u></u>	
16	Sediment lost during recovery. — End of Core —	End of core	End of core	End of core	End of core
18		-			

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 5-3 Rep 3

Maximum depth of retained sediment: #DIV/0! ft

Mudline elevation: -14.2 ft (Corps lake datum)

	Core	Laboratory	
	collection	processing	
Date:	8/10/2004	Aug. 10, 2004	
Time:	12:40	16:00	

Percent recovery (on-deck): 70%

0 -	Visual Description of Sediment	Summary Interpretation	Segment End of core ———	Primary Sample ID	Secondary Sample ID End of core
•	-	+			
	- -				
-	-	+		34011008008	
1 -					
-	-	1			
	-	+			
1 -		+			
-	Black; silty clay with sand and wood fragments; Very soft, wet. Black oil	- Upper recent lake			
-	blebs and strong asphalt odor.	deposits			
2 -	+	+			
	-	+			
~		+			
Depth below mudline (ft.)	•	+			
alline .	-				
, mu		-			
≥ 3 -					
pel -	-	Ì			
epth	Dark olive grading down to dark gray;	+		34011008009	
<u>ŏ</u> 3-	organic clay (OL) and wood pieces	 – – Lower recent lake - – – deposits 		34011008003	
-	or oil stains.	+			
-	-	+			
4 -					
	-	+			
-					
4 -	Dark gray sandy gravel (GW); Dense, moist to wet. No oil sheen or stains.	Stratified Drift			
-	-	+			
-	-	-		34011008010	
5 -	Sediment Ic End of Core e recovery.	End of core		End of core	
-	-	ł			
-	+	Į			
5 -		L			

Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

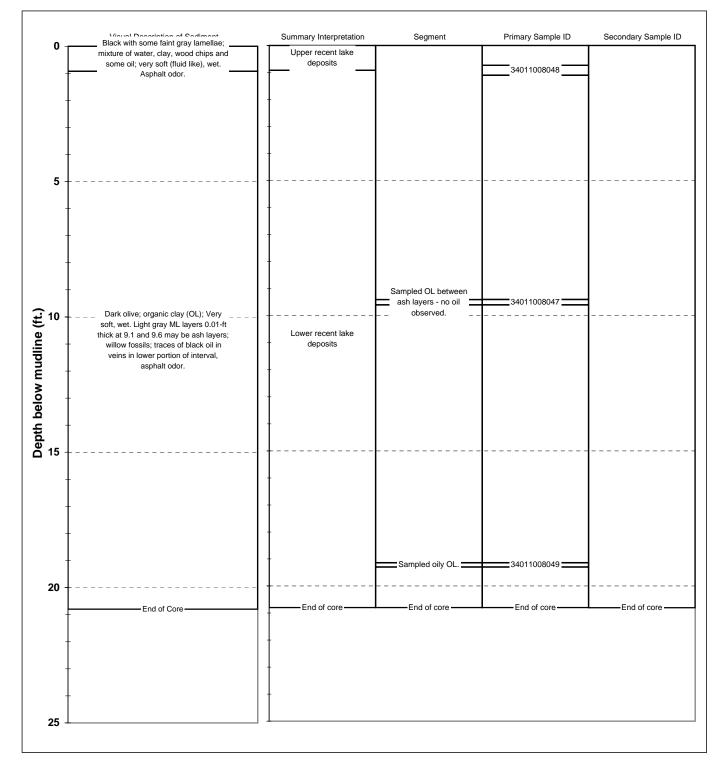
Station: 5--2

Maximum depth of retained sediment: 20.8 ft

Mudline elevation: -20.1 ft (Corps lake datum)

	Core Laboratory	
	collection	processing
Date:	8/12/2004	Aug. 12, 2004
Time:	11:30	15:45

Percent recovery (on-deck): 78%



Project: North Lake Union Sediment Survey Project No: COS-NAPL.T5

Station: 5--4

Maximum depth of retained sediment: 13.9 ft

Mudline elevation: -19.1 ft (Corps lake datum)

	Core Laborato	
	collection	processing
Date:	8/12/2004	Aug. 13, 2004
Time:	13:15	10:30

ſ

Percent recovery (on-deck): 76%

Field Log: J. LaManna

Summary Log: J. LaManna

		Visual Description of Sediment	Summary Interpretation	Segment	Primary Sample ID	Secondary Sample ID
	°]		- Upper recent lake			
]	Gray grading down to dark gray or black; clay (CL) with wood fragments	deposits		34011008051	
		and blebs and streaks of black tarry oil; Very soft, wet. Asphalt or creosote	Ţ			
	-	odor.				
	2		-			
	-	-	+			
	-	-	+			
	1	-	-			
			+			
	4		1			
			I			
		-				
	-	-				
_	6		+			
(£f.)	-	-	-			
ne	1	Dark olive gray; organic clay (OL) with	Lower recent lake			
ilbu	1	traces of roots; very soft, wet. Thin bed of oily, f-m sand (SP) at 14.5. Veins	_ deposits			
Ē	8	with black tarry oil throughout interval	T			
Depth below mudline (ft.)	Ĭ	_	I			
þe	-					
pth	-	-	-			
	-	-	+			
	10 -		Ŧ			
			-			
	1		-			
		_	I			
	12					
	-	-				
	-	-	+	Sampled oily SP.	34011008050	
	-	-	-			
		-	End of core	End of core	End of core	End of core
	14 -	End of Core	+			
	1	-	1			
]		Į			
		-				
	I6]					

SUB-ATTACHMENT 2D-4.3 RI/FS 2005 Sediment Core Logs Seattle Law Department Gas Works Park

Chemical Forensics Sampling Report

Appendix C RI/FS 2005 Sediment Core Logs

DRAFT

CONFIDENTIAL AND PRIVILEGED: PREPARED IN PREPARATION FOR LITIGATION

Project: Gas Works Sediment-Western Study Area Project No: 3400542.002

Station: GWS-EC01

Maximum depth of retained sediment: 20.1 ft

Mudline elevation: -19.9 ft (Corps lake datum)

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Percent recovery (on-deck): 76%

	Core	Laboratory				
	collection	processing	Position:	N238271	E1269512	(NAD83 SPC WAN)
Date:	5/18/2005	5/18/2005	Field Log:	John LaManna		
Time:	13:01	0:00	Summary Log:	John LaManna		

0 _Ŧ	Visual Description of Sediment	Summary Interpretation	Symbol	Primary Sample ID	Secondary Sample
1	Thinnly bedded to laminated brown organic silt and gray clay, with some wood fragments and fibers. Very soft; wet.	Upper recent lake - deposits	OH/CI		
2					
3		ŧ		GWS-EC01-0025	
4		ŧ			
5		ŧ			
6		I I I I I I I I I I I I I I I I I I I			
ŧ					
7		<u>+</u>			
8					
9		<u>+</u>			
10 1		ŧ		GWS-EC01-0090	
11	Dark brown, organic silt with sand; amorphous with trace plant fibers;	Lower recent lake deposits	OH		
12	massive. Very soft; moist to wet.				
ŧ					
13		+			
14		I			
15		ŧ			
16 1		<u></u>			
17 1					
ŧ					
18		<u>+</u>		GWS-EC01-0172	
19		ŧ			
		Ŧ			1

Project: Gas Works Sediment-Western Study Area Project No: 3400542.002

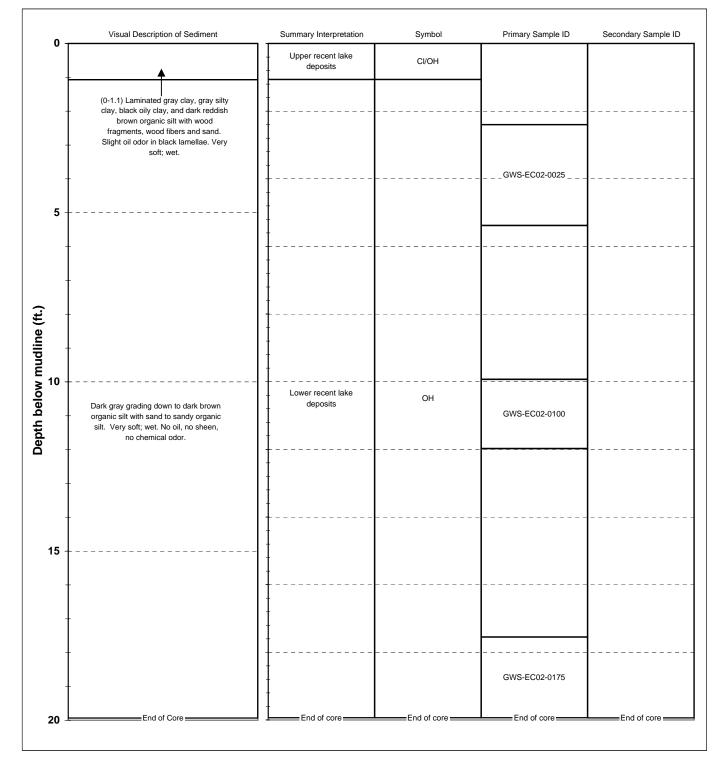
Station: GWS-EC02

Maximum depth of retained sediment: 19.9 ft

Mudline elevation: -20.3 ft (Corps lake datum)

Percent recovery (on-deck): 61%

	Core	Laboratory				
	collection	processing	Position:	N238520	E1269647	(NAD83 SPC WAN)
Date:	5/19/2005	5/19/2005	Field Log:	John LaManna		
Time:	8:41	0:00	Summary Log:	John LaManna		



Project: Gas Works Sediment-Western Study Area Project No: 3400542.002

Station: GWS-EC03

Maximum depth of retained sediment: 20.3 ft

Mudline elevation: -19.9 ft (Corps lake datum)

Percent recovery (on-deck): 70%

E1269515

	Core	Laboratory	
	collection	processing	Pos
Date:	5/19/2005	5/19/2005	Field
Time:	10:28	0:00	Summary

sition: N238690 d Log: John LaManna y Log: John LaManna

(NAD83 SPC WAN)

0 -	Visual Description of Sediment	Summary Interpretation	Symbol	Primary Sample ID	Secondary Sample ID
F	↑	Upper recent lake deposits	OH/CI	-	
+	(0-0.9) Interbedded and laminated; dark reddish brown, organic silt and gray, dark gray and black, silty clay with few wood fibers and sand. Black oil stains and oil odor. Very soft; wet.			GWS-EC03-0015	
5 -				GWS-EC03-0055	
÷ 10 +	Dark gray grading down to dark brown, sandy organic silt to organic silt with			GWS-EC03-0078	
	sandy organic sin to organic sin with a sandy organic sin with a sandy organic sin to organic sin with a sandy but smell decreases with depth. Bottom smells oily. Black oil veinlets at 10.7-ft.	Lower recent lake deposits	ОН	GWS-EC03-0110	
15 -				GWS-EC03-0137	
				GWS-EC03-0173	
20 -	End of Core	End of core	End of core ———	End of core	End of core —
25		-			

Project: Gas Works Sediment-Western Study Area Project No: 3400542.002

Station: GWS-EC04

Maximum depth of retained sediment: 6.9 ft

Mudline elevation: -19.1 ft (Corps lake datum)

Percent recovery (on-deck): 51%

	Core	Laboratory				
	collection	processing	Position:	N238646	E1269721	(NAD83 SPC WAN)
Date:	5/17/2005	5/17/2005	Field Log:	John LaManna		
Time:	10:30	0:00	Summary Log:	John LaManna		

0	Visual Description of Sediment	Summary Interpretation	Symbol	Primary Sample ID	Secondary Sample ID
1	Black clay with undecayed wood fragments; trace oil sheen. Very soft, wet.	Upper recent lake deposits	C!?		
2	Gray clay with silt. Very soft; wet. No oil.	Upper recent lake	Cl		
3 3 4 4	Dark brown, sandy organic silt to organic silt with sand. Very soft; wet. Slight asphalt odor.	Lower recent lake deposits	Pt		
		- - - - - -		GW3-EU04-0023	
6	Interbedded dark brown, sandy organic silt and poorly graded sand. Some medium and coarse angular sand grains, few quartz sand grains Trace black oil in sand bed; asphaltic odor. Igneous rock stuck in catcher suggests refusal in stratified drift.	Lower recent lake	Pt/Sp		
7 + -	End of Core	End of core	End of core	End of core	End of core
8		-			

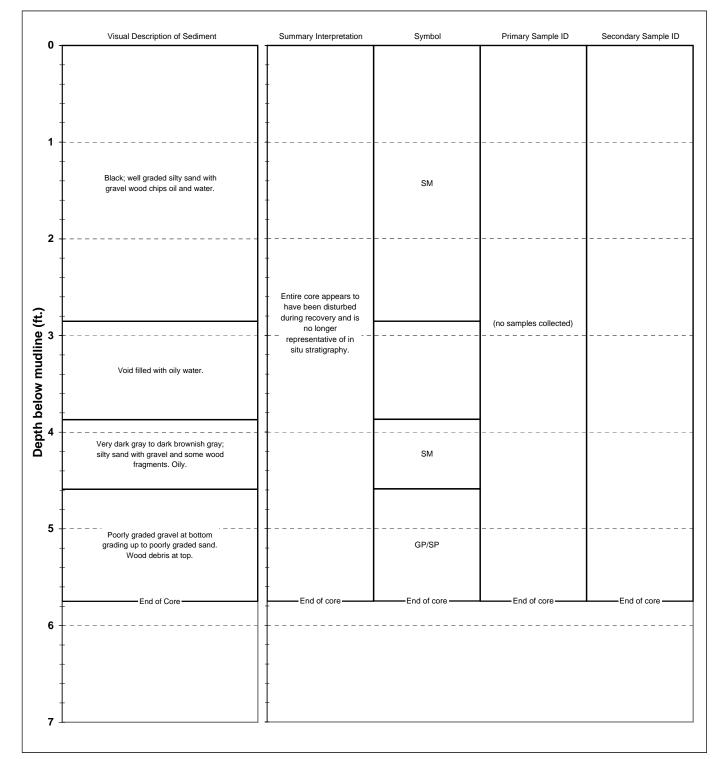
Project: Gas Works Sediment-Western Study Area 3400542.002 **Project No:**

Station: GWS-EC05

Maximum depth of retained sediment: ft 5.7

Mudline elevation: -7.7 ft (Corps lake datum) Percent recovery (on-deck): 75%

	Core	Laboratory				
	collection	processing	Position:	N238693	E1269888	(NAD83 SPC WAN)
Date:	5/16/2005	5/16/2005	Field Log:	John LaManna		
Time:	13:10	0:00	Summary Log:	John LaManna		



Project: Gas Works Sediment-Western Study Area **Project No:** 3400542.002

Station: GWS-EC05R4

Maximum depth of retained sediment: 7.7 ft

Mudline elevation: -10.2 ft (Corps lake datum)

Percent recovery (on-deck):

58%

	Core	Laboratory				
	collection	processing	Position:	N238680	E1269868	(NAD83 SPC WAN)
Date:	5/20/2005	5/20/2005	Field Log:	John LaManna		
Time:	12:50	0:00	Summary Log:	John LaManna		

0 T	Visual Description of Sediment	Summary Interpretation	Symbol	Primary Sample ID	Secondary Sample ID
-	Dark gray slurry of water, clay, silt, wood chips and oil. Asphalt odor.	-	CI?		
1 -					
-	- - Black mixture of clay, water, oil, wood	- Upper recent lake - deposits -	Cl?		
2 -	fragments and sand. Gelatinous - - -				
3 -					
ine (ft.)	Dark reddish brown, organic silt. Very soft; wet. Gelatinous.	Lower recent lake deposits	он	(no samples collected)	
low mud	- Gray; silty gravel. Very loose; wet.	Weathered stratified	GM		
Depth below mudline (ft.)	Gray; sandy silt with gravel and some — — — — interbeds of silty fine sand. Dense; moist to wet. No oil.		SM		
6 -	- Gray; poorly graded fine sand, <20% gravel, rounded, <5% silt. Dense. No oil.	 Stratified drift: possibly glacially overridden. 	SP		
7 -	- - - – – – Gray; silty fine sand. Dense; moist to– – – – - wet. No oil.		SM		
8 -	End of Core	End of core	End of core ———	End of core	End of core ——
-	-	-			
9		1			

Project: Gas Works Sediment-Western Study Area Project No: 3400542.002

Station: GWS-EC06

Maximum depth of retained sediment: 8.1 ft

Mudline elevation: -11.8 ft (Corps lake datum)

Percent recovery (on-deck): 71%

	Core	Laboratory				
	collection	processing	Position:	N238765	E1269800	(NAD83 SPC WAN)
Date:	5/16/2005	5/16/2005	Field Log:	John LaManna		
Time:	11:16	0:00	Summary Log:	John LaManna		

0 —	Visual Description of Sediment	Summary Interpretation	Symbol	Primary Sample ID	Secondary Sample ID
1+					
2 +	Black grading down to gray; mixture of water, clay, wood fragments, and oil. Laminated clay over abundant wood chips in bottom foot. Very soft; wet. Blebs of black oil. Asphalt odor.	- Upper recent lake deposits	CI?		
3 +		- 		GWS-EC06-0023	
4		+ - - - -		GWS-EC06-0038	
5 + - - 6 +	Dark gray and black; poorly graded sandy silt; with black oil from 4.5 ft to 4.7 ft and 6.5 ft to 6.7 ft (on top of clay); one oil-saturated undecayed wood fragment 0.3-ft long. Loose; wet.		SP	 GWS-EC06-0056	
7 +	Gray; lean clay overlying gray, poorly graded sand with gravel and a rounded cobble. Cobble is granodiorite.		CL/GW		
8 +	End of Core	End of core	End of core	End of core	End of core
9		-			

Project: Gas Works Sediment-Western Study Area **Project No:** 3400542.002

Station: GWS-EC07

Maximum depth of retained sediment: 7.7 ft

Mudline elevation: -2.5 ft (Corps lake datum)

Percent recovery (on-deck): 84%

	Core	Laboratory				
	collection	processing	Position:	N239035	E1269642	(NAD83 SPC WAN)
Date:	5/16/2005	5/16/2005	Field Log:	John LaManna		
Time:	10:07	0:00	Summary Log:	John LaManna		

• ⊤	Visual Description of Sediment	Summary Interpretation	Symbol	Primary Sample ID	Secondary Sample I
+ + + + 1 +	Black; emulsion of water, wood fragments, oil and sand. Gelatinous to semi-solid. Very soft; wet. Asphalt odor.	- Upper recent lake deposit	(NA)		
2 + -	Very dark olive brown; silty sand. Trace wood fragments, possible brick fragments. Thin bed of black grit at 10.6. Oily. Very loose; wet.		GW	GWS-EC07-0013	
3	Black; silty sand, at bottom of interval, fine to coase gravel; trace wood	Fill or disturbed lake	OL?	GWS-EC07-0034	
4	fragments and glass. Blebs of black oil.			GWS-EC07-0046	
6	Gray; beds of clay, over poorly graded sand, over gravel with sand, over silty sand. Medium dense; wet. Some oil sheen on sand bed.		CL/SP/GW/SM	GWS-EC07-0056	
7 -		- - - - - -			
8	End of Core	End of core	End of core ———	End of core	End of core
‡ و					

Project: Gas Works Sediment-Western Study Area Project No: 3400542.002

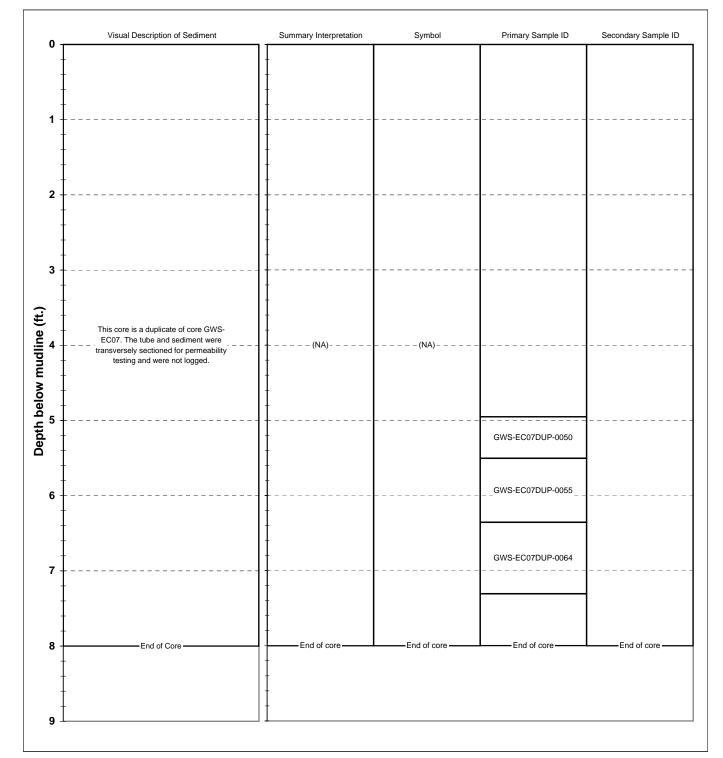
Station: GWS-EC07DUP

Maximum depth of retained sediment: 8.0 ft

Mudline elevation: -1.6 ft (Corps lake datum)

Percent recovery (on-deck): 78%

	Core	Laboratory				
	collection	processing	Position:	N239033	E1269643	(NAD83 SPC WAN)
Date:	5/16/2005	5/16/2005	Field Log:	John LaManna		
Time:	14:36	0:00	Summary Log:	John LaManna		



Project: Gas Works Sediment-Western Study Area **Project No:** 3400542.002

Station: GWS-EC08

Maximum depth of retained sediment: 15.6 ft

Mudline elevation: -19.8 ft (Corps lake datum) 85%

Percent recovery (on-deck):

	Core	Laboratory	
	collection	processing	
Date:	5/16/2005	5/16/2005	
Time:	13:54	0:00	Sum

Position: N238894 Field Log: John LaManna mary Log: John LaManna

E1269444

(NAD83 SPC WAN)

0 	Visual Description of Sediment	Summary Interpretation	Symbol	Primary Sample ID	Secondary Sample ID
	•	- Upper recent lake deposits	OH/CL		
2	(0-1.2) Laminated, dark brown; organic silt and very dark gray silty clay, with wood fragments and sand overlying black grading down to gray, silty clay, with wood fragments. Oily. Very soft; wet.			GWS-GC08-0006	
4 + -		- 		GWS-GC08-0028	
6 + -					
-				GWS-GC08-0048	
uiue 8 8		Lower recent lake			
Leptn below mualine (π)	Grayish brown grading down to yellowish brown;organic silt with sand to sandy organic silt, trace plant fibers. Very soft; moist to wet. Moisture content decreases with depth. Smells	deposits	ОН	GWS-GC08-0068	
10 + - 	$-$ - $-$ oily. Oil veinlets at 6.3 ft and 8.7 ft. \cdot	+		GWS-GC08-0100	
12 + -					
14 -					
ţ				GWS-GC08-0129	
16 - -	End of Core ————————————————————————————————————	End of core	End of core	End of core	End of core ———
18		-			

Project: Gas Works Sediment-Western Study Area Project No: 3400542.002

Station: GWS-EC09

Maximum depth of retained sediment: 20.3 ft Percent recovery (on-deck): 76%

E1269353

Mudline elevation: -19.6 ft (Corps lake datum)

	Core	Laborator
	collection	processing
Date:	5/19/2005	5/19/2005
Time:	11:06	0:00

ooratory cessing 9/2005 0:00 S

Position: N238842 Field Log: John LaManna Summary Log: John LaManna

(NAD83 SPC WAN)

Visual Description of Sediment Primary Sample ID Secondary Sample ID Summary Interpretation Symbol 0 (0-0.15) Upper recent (0-0.15) Laminated, gray clay and dark (0-0.15) CL/OH reddish brown organic silt; Very soft; lake deposits wet. GWS-GC09-0008 5 Gray grading down to dark brown; Depth below mudline (ft.) 10 organic silt with sand to sandy organic _ GWS-GC09-0092 Lower recent lake silt, massive, moisture content OH deposits deceases with depth. Top 0.1-ft is gray. Very soft; wet. GWS-GC09-0176 20 End of core End of core End of core End of core --End of Core-

25

Project: Gas Works Sediment-Western Study Area **Project No:** 3400542.002

Station: GWS-EC09DUP/EC24

Maximum depth of retained sediment: 20.3 ft Percent recovery (on-deck):

Mudline elevation: -19.7 ft (Corps lake datum)

	Core collection	Laborato processi
Date:	5/20/2005	5/20/200
Time:	10:35	0:00

ory ing 05

Position: N238845 Field Log: John LaManna Summary Log: John LaManna

E1269353

(NAD83 SPC WAN)

0 —	Visual Description of Sediment	Summary Interpretation	Symbol	Primary Sample ID	Secondary Sample
-	↑		^		
	I (0-0.45) Organic silt, clay, wood fragments, water and oil (asphalt odor) overlying laminated, dark brown organic silt and gray clay with silt. Very soft; wet.	Upper recent lake deposits	I OH/CL	GWS-EC-0008	
5 +					
10	Gray grading down to dark brown, sandy organic silt to organic silt with sand, amorphous, massive, water content decreases with depth, trace visible plant parts. Gray color in top 0.3 ft. Very soft; wet. No odor in most of this interval; no odor at bottom	Lower recent lake deposits	он	GWS-EC-0092	
15					
-		-		GWS-EC-0176	
20 +	End of Core	End of core	End of core	End of core	End of core
ļ		+			

77%

Project: Gas Works Sediment-Western Study Area 3400542.002 **Project No:**

Station: GWS-EC10

Maximum depth of retained sediment: 20.3 ft Percent recovery (on-deck): 82%

Mudline elevation: -19.5 ft (Corps lake datum)

	Core	Laborator
	collection	processing
Date:	5/18/2005	5/18/2005
Time:	13:55	0:00

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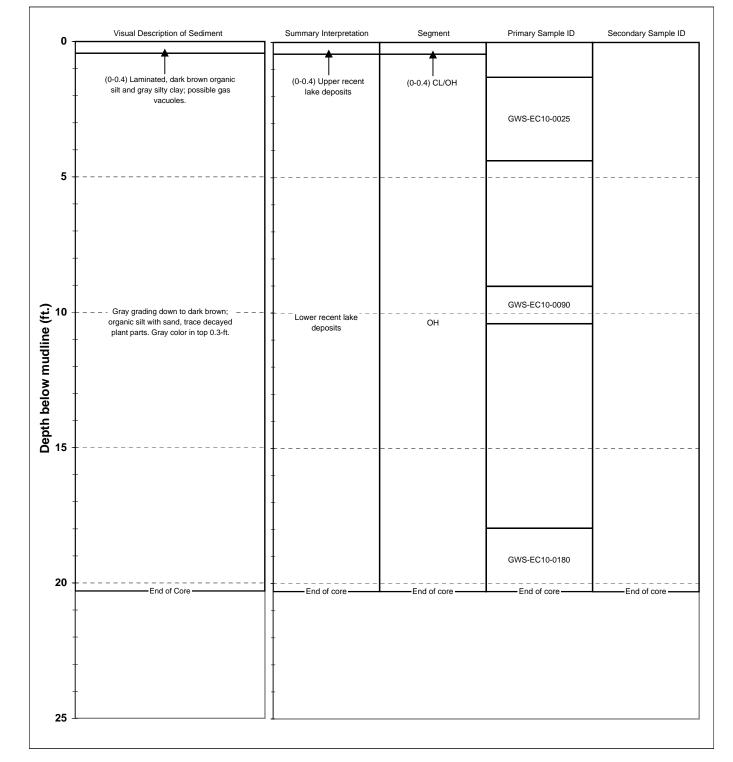
N238540 Field Log: John LaManna

(NAD83 SPC WAN)

Position: Summary Log:

John LaManna

E1269116



Project: Gas Works Sediment-Western Study Area **Project No:** 3400542.002

Station: GWS-EC11

Maximum depth of retained sediment: 8.8 ft

Mudline elevation: 19.3 ft (Corps lake datum)

Γ

Percent recovery (on-deck): 69%

	Core	Laboratory				
	collection	processing	Position:	N239263	E1269530	(NAD83 SPC WAN)
Date:	5/16/2005	5/16/2005	Field Log:	John LaManna		
Time:	9:15	0:00	Summary Log:	John LaManna		

0 Τ	Visual Description of Sediment	Summary Interpretation	Symbol	Primary Sample ID	Secondary Sample ID
- - - - - - - - - - - - - - - - - - -	Dark gray poorly graded sand with gravel overlying dark gray silty sand – – – with gravel; trace porceline, wood and – – – glass fragments. No chemical odor.	- - - - Fill - -	SP/SM		
2 -		- - 			
3 -	- - - - -	- - - - 		GWS-EC11-0019	
4	- - - 	- - - - - -			
4	Dark grayish brown, poorly-graded, fine sand with trace rounded gravel and trace coarse sand, a lump of organics or decayed wood; fine gravel size lump of tarry sand with asphalt odor.	- - - Fill 	SP	GWS-EC11-0048	
6 -	- 	- 			
7	- - 	 			
8		- - 		GWS-EC11-0067	
	Dark brown silt; over 0.1 ft of dark brown wood fragments. End of Core	Fill End of core	ML End of core	End of core	End of core
9 -		+ - -			
10	-	Ŧ			

Project: Gas Works Sediment-Western Study Area **Project No:** 3400542.002

Station: GWS-EC11Dup/EC-23

Maximum depth of retained sediment: 7.8 ft

Mudline elevation: 19.5 ft (Corps lake datum)

Percent recovery (on-deck): 68%

	Core collection	Laboratory processing	Position:	N239262	E1269530	(NAD83 SPC WAN)
Date: Time:	5/17/2005 12:56	5/17/2005 0:00	Field Log: Summary Log:	John LaManna John LaManna		· · · ·

0	Visual Description of Sediment	Summary Interpretation	Symbol	Primary Sample ID	Secondary Sample ID
	Gray; poorly graded fine sand. No chemical odor, no oil.	- - Fill	SP		
1 - - - 2	Gray; poorly graded fine sand with 15- 20% rounded and angular gravel, with glass and with undecayed wood. No oil, no chemical odor.	- - - Fill -	SP		
3 + -				GWS-EC23-0019	
	Gray, with strong brown mottles toward				
	bottom of interval; poorly graded fine sand with trace coarse gravel and a dark brown silt lump. More oxidized in bottom 0.5-ft.	- Fill - Fill 	SP	GWS-EC23-0048	
6 + -					
7 + -	(7.4-7.8) Laminated, brown silty fine sand and sandy silt, grading down to gravel with silt and wood chips.	- 		GWS-EC23-0067	
8 + -	End of Core	- Fill End of core	SM? or GM?	End of core	End of core
9					

Project: Gas Works Sediment-Western Study Area Project No: 3400542.002

Station: GWS-EC12

Maximum depth of retained sediment: 15.5 ft Percent recovery (on-deck): 73%

Mudline elevation: -19.0 ft (Corps lake datum)

	Core collection	Laboratory processing
Date:	5/17/2005	5/17/2005
Time:	8:46	0:00

Position:N239177Field Log:John LaMannaSummary Log:John LaManna

E1269370 (NAD83

(NAD83 SPC WAN)

•					
0	_	Upper recent lake deposits	CI		
Į		I I			
-	(0-0.7 ft) Black to gray emulsion of water, oil, silt and clay. Gray clay at				
2 +	- bottom of interval. Very soft; wet	•			
ł	-	-		GWS-EC12-0008	
ł	-	+			
ł	-	+			
4		+			
ł	-	+			
ļ	-				
ł	-	+		GWS-EC12-0043	
6 +		+			
t	Dark gray and black organic silt with	†			
Ţ	sand and oil grading to dark brown organic silt with sand, amorphous,	Lower recent lake	ОН		
ļ	massive, trace decayed plant parts.	deposits		GWS-EC12-0064	
8 -	Very soft; moist to wet. No oil below a	+			
ł	-	+ I			
ł	-	+ I			
ŧ	-	†		GWS-EC12-0084	
, †		†			
10 +		Ţ			
ļ	-	1			
ļ	-	+ I			
ł	-	+			
12 +		+			
ł	-	†			
t	-	†			
ļ				GWS-EC12-0108	
14]	Dark brown sandy organic silt	L			
·-	graded fine and medium sand. Trace	Lower recent lake	011/05		
ļ	angular (with sharp edges), tabular, coarse sand-sized rock or shell	deposits	OH/SP		
ł	fragments with sand. Oil in sand beds.	+ I			
ŧ	End of Core	End of core	End of core	End of core	End of core
16 +		+			
ļ	-	† +			
ļ	-	ţ			
18 [⊥]					

Project: Gas Works Sediment-Western Study Area 3400542.002 Project No:

Station: GWS-EC13

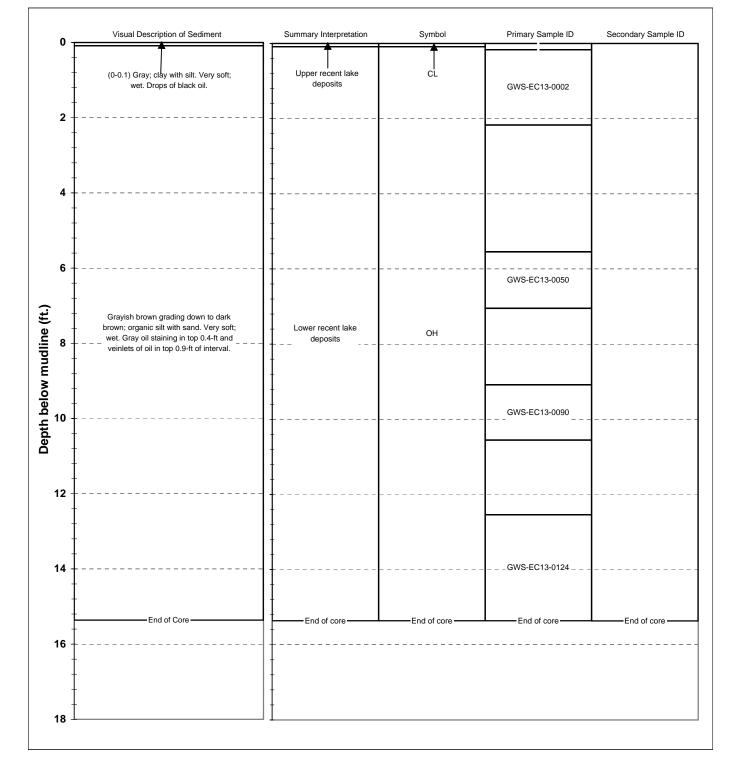
Maximum depth of retained sediment: 15.4 ft Percent recovery (on-deck): 73%

Mudline elevation: -20.4 ft (Corps lake datum)

	Core collection	Laboratory processing
Date:	5/17/2005	5/17/2005
Time:	9:39	0:00

Position: N239061 Field Log: John LaManna Summary Log: John LaManna

E1269278 (NAD83 SPC WAN)



Project: Gas Works Sediment-Western Study Area Project No: 3400542.002

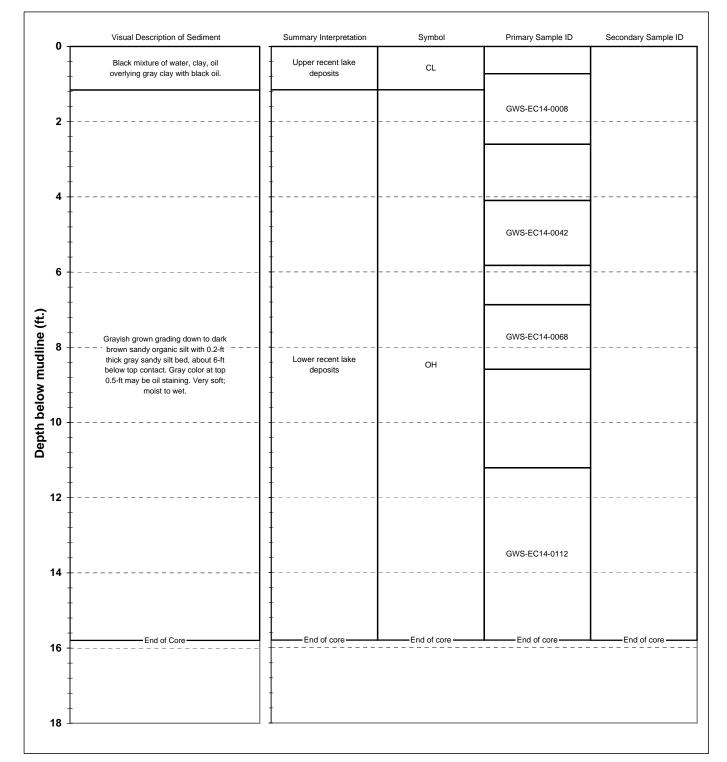
Station: GWS-EC14

Maximum depth of retained sediment: 15.8 ft

Mudline elevation: -20.5 ft (Corps lake datum)

num deput of retained sediment.	10.0
Percent recovery (on-deck):	71%

	Core collection	Laboratory processing	Position:	N239240	E1269295	(NAD83 SPC WAN)
Date:	5/17/2005	5/17/2005	Field Log:	John LaManna		· · · · · · · · · · · · · · · · · · ·
Time:	13:35	0:00	Summary Log:			



Project: Gas Works Sediment-Western Study Area 3400542.002 **Project No:**

Station: GWS-EC15

Maximum depth of retained sediment: 19.6 ft Percent recovery (on-deck): 71%

Primary Sample ID

Mudline elevation: -19.3 ft (Corps lake datum)

0

	Core	Laborato
	collection	processin
Date:	5/20/2005	5/20/2005
Time:	9:30	0:00

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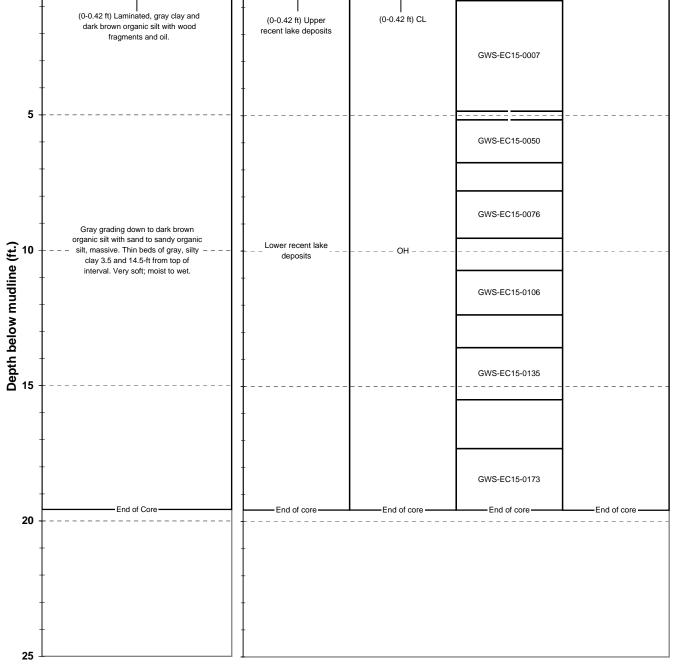
Position: N239001 John LaManna

E1269161

(NAD83 SPC WAN)

Secondary Sample ID

)5 Field Log: John LaManna Summary Log: Visual Description of Sediment Summary Interpretation Symbol (0-0.42 ft) Laminated, gray clay and (0-0.42 ft) Upper dark brown organic silt with wood recent lake deposits fragments and oil.



Project: Gas Works Sediment-Western Study Area Project No: 3400542.002

Station: GWS-EC16

Maximum depth of retained sediment: 20.0 ft

Mudline elevation: -19.0 ft (Corps lake datum)

Percent recovery (on-deck): 74%

(NAD83 SPC WAN)

	Core	Laboratory			
	collection	processing	Position:	N238874	E1268806
Date:	5/18/2005	5/18/2005	Field Log:	John LaManna	
Time:	10:58	0:00	Summary Log:	John LaManna	

Visual Description of Sediment Symbol Primary Sample ID Secondary Sample ID Summary Interpretation 0 Upper recent lake deposits CL Gray clay. Very soft; wet. GWC-EC16-0010 5 GWC-EC16-0066 Depth below mudline (ft.) 10 Dark brown organic silt with sand. Very soft; moist to wet. Lower recent lake OH deposits GWC-EC16-0173 20 End of core End of core End of core-End of core End of Core 25

Project: Gas Works Sediment-Western Study Area **Project No:** 3400542.002

Station: GWS-EC17

Maximum depth of retained sediment: 20.1 ft Percent recovery (on-deck): 74%

	Core	Laboratory
	collection	processing
Date:	5/20/2005	5/20/2005
Time:	8:25	0:00

Position:

N238760 Field Log: John LaManna Summary Log: John LaManna

E1269052 (NAD83 SPC WAN)

0 -	Visual Description of Sediment	Summary Interpretation	Symbol	Primary Sample ID	Secondary Sample ID
+	(0-0.44 ft) Dark brownish gray organic clay with silt. Trace wood; no oil, sheen or chemical odor.	(0-0.44 ft) Upper recent lake deposits	↑ (0-0.44 ft) OL	GWS-EC17-0009	
5 -					
-		-		GWS-EC17-0055	
+ 10 + + + + + + + + + + + + + + + + + +	Dark grayish brown grading down to dark brown; organic silt with sand. Grayish color in upper 0.3-ft. Very soft; wet. No oil, sheen or chemical odor.	Lower recent lake deposits -	он		
		-		GWS-EC17-0125	
-		-			
20 -	End of Core		End of core	GWS-EC17-0178	End of core
+		-			
25		-			

Project: Gas Works Sediment-Western Study Area Project No: 3400542.002

Station: GWS-EC18

Maximum depth of retained sediment: 15.6 ft

Mudline elevation: -19.5 ft (Corps lake datum)

Percent recovery (on-deck): 70%

	Core	Laboratory		
	collection	processing	Position:	N239112
Date:	5/20/2005	5/20/2005	Field Log:	John LaManna
Time:	11:15	0:00	Summary Log:	John LaManna

E1268852 (NAD83 SPC WAN)

Visual Description of Sediment Primary Sample ID Secondary Sample ID Summary Interpretation Symbol 0 Upper recent lake OH/CL deposits (0-0.8 ft) Laminated, dark brown 2 organic silt and gray clay with silt. Very soft; moist. GWS-EC18-0022 4 6 Depth below mudline (ft.) GWS-EC18-0068 Dark gray grading down to dark brown, 8 organic silt with sand, amorphous, Lower recent lake OH massive. Gray color in top 0.5-ft of deposits interval. Very soft; wet. 10 12 GWS-EC18-0127 14 End of Core End of core End of core -End of core -End of core 16 18

Project: Gas Works Sediment-Western Study Area 3400542.002 **Project No:**

Station: GWS-EC19

Maximum depth of retained sediment: 20.3 ft Percent recovery (on-deck): 79%

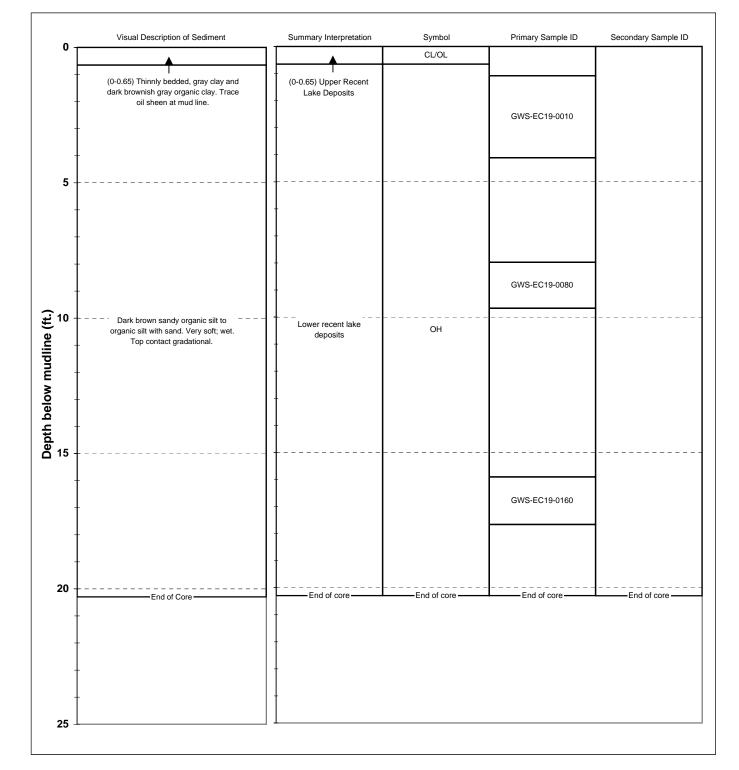
Mudline elevation: -19.1 ft (Corps lake datum)

	Core	Laborator
	collection	processin
Date:	5/18/2005	5/18/2005
Time:	9:07	0:00

·у g 5

Position: N239034 Field Log: John LaManna Summary Log: John LaManna

E1268729 (NAD83 SPC WAN)



Project: Gas Works Sediment-Western Study Area **Project No:** 3400542.002

Station: GWS-EC20

Maximum depth of retained sediment: 13.9 ft

E1269361

Mudline elevation: -19.5 ft (Corps lake datum)

Percent recovery (on-deck):

69%

	Core	Laboratory	
	collection	processing	
Date:	5/19/2005	5/19/2005	
Time:	9:38	0:00	

Position: N238479

Field Log: John LaManna Summary Log: John LaManna (NAD83 SPC WAN)

0 -	Visual Description of Sediment	Summary Interpretation	Symbol	Primary Sample ID	Secondary Sample ID
	†	Upper recent lake deposits	CL/OH		
2 -	I (0-0.8 ft) Laminated, dark red-brown organic silt and gray to black clay(?). Trace wood fibers. Very soft; wet.	-			
+				GWC-EC20-0018	
4 +					
6		-			
	Gray grading down to dark brown organic silt with sand. Gray color and	- Lower recent lake	ОН	GWC-EC20-0058	
8 -	trace gas vacuoles in top 0.5-ft. Very soft; moist to wet.	deposits			
8 +					
10 +		+			
		-			
12 +				GWC-EC20-0128	
14 -	End of Core	End of core	End of core	End of core	End of core
16		1			

Project: Gas Works Sediment-Western Study Area Project No: 3400542.002

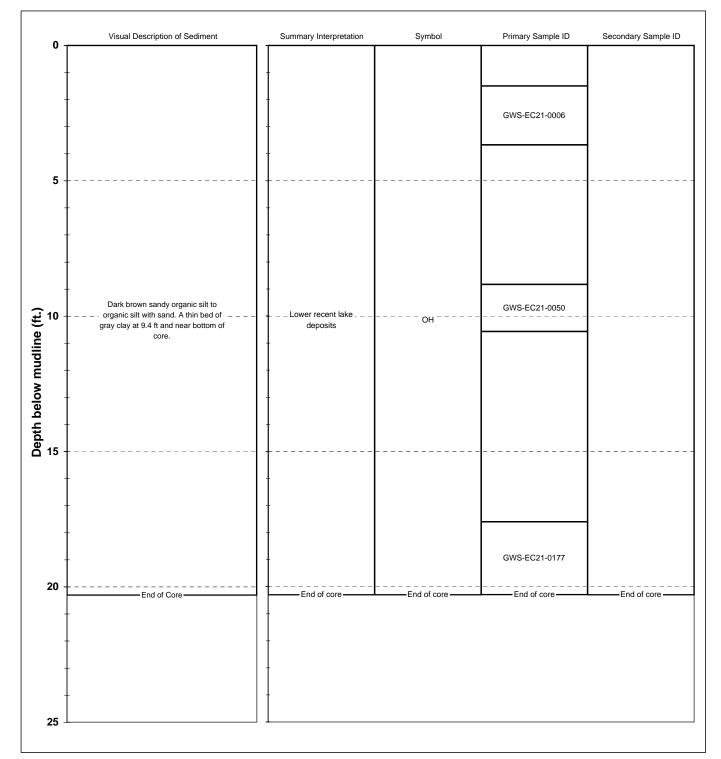
Station: GWS-EC21

Maximum depth of retained sediment: 20.3 ft

Mudline elevation: -18.8 ft (Corps lake datum)

Percent recovery (on-deck): 77%

	Core	Laboratory				
	collection	processing	Position:	N238265	E1269126	(NAD83 SPC WAN)
Date:	5/18/2005	5/18/2005	Field Log:	John LaManna		
Time:	12:10	0:00	Summary Log:	John LaManna		



Project: Gas Works Sediment-Western Study Area **Project No:** 3400542.002

Station: GWS-EC22

Maximum depth of retained sediment: 19.8 ft Percent recovery (on-deck): 73%

Mudline elevation: -17.8 ft (Corps lake datum)

	Core	Laboratory
	collection	processing
Date:	5/17/2005	5/17/2005
Time:	14:25	0:00

Position: N238694 Field Log: John LaManna Summary Log: John LaManna

E1268638 (NAD83 SPC WAN)

Visual Description of Sediment Summary Interpretation Symbol Secondary Sample ID Primary Sample ID 0 CL (0-0.7) Upper recent (0-0.7) Gray, clay with silt. Very soft; lake deposits wet. GWS-EC22-0015 5 GWS-EC22-0055 GWS-EC22-0078 Dark brown, sandy organic silt to Depth below mudline (ft.) 21 01 organic silt, moisture content Lower recent lake OH decreases with depth. Very soft; moist deposits. to wet. Top contact gradational. GWS-EC22-0105 GWS-EC22-0135 GWS-EC22-0162 _____End of Core End of core -End of core End of core -End of core-20 ____ 25

SUB-ATTACHMENT 2D-4.4 Photographs of NAPL Investigation 2004 Sediment Cores Seattle Law Department Gas Works Park

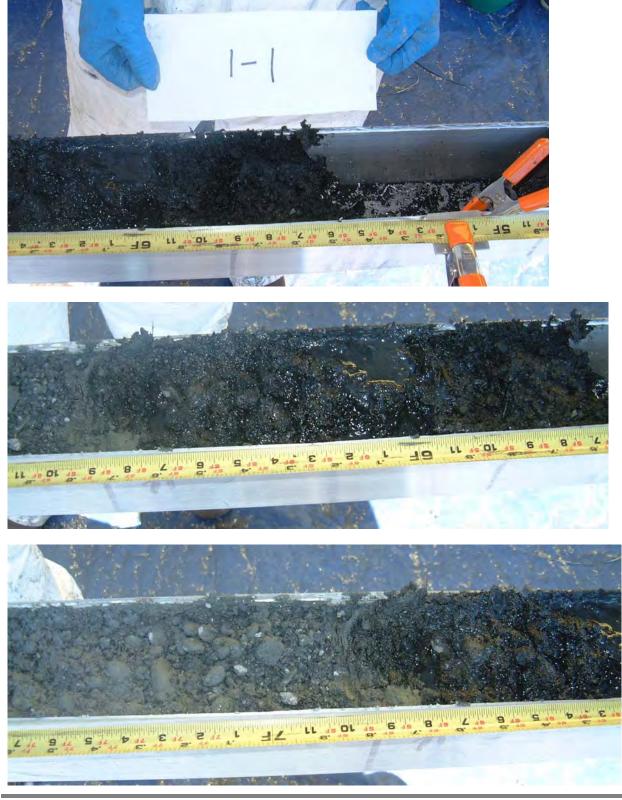
Chemical Forensics Sampling Report

Appendix D Photographs of NAPL Investigation 2004 Sediment Cores

DRAFT

CONFIDENTIAL AND PRIVILEGED: PREPARED IN PREPARATION FOR LITIGATION

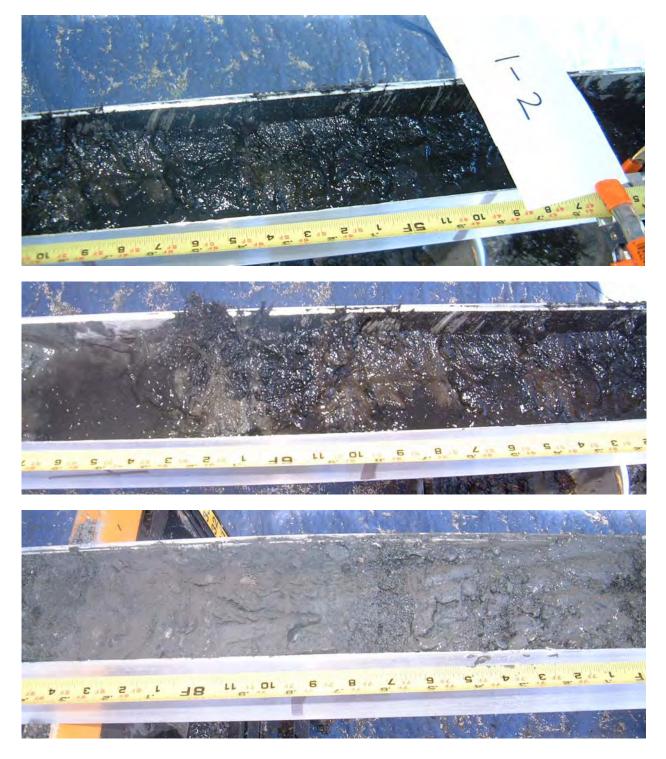
Photographs of NAPL Investigation Core 1-1: Starting at the top of the core

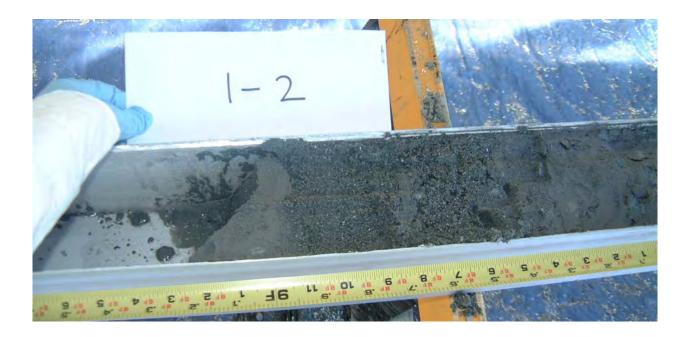




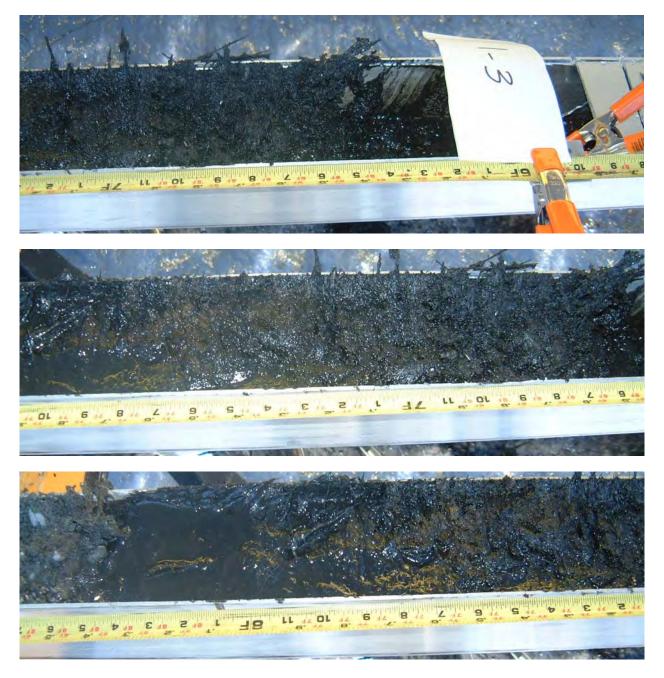


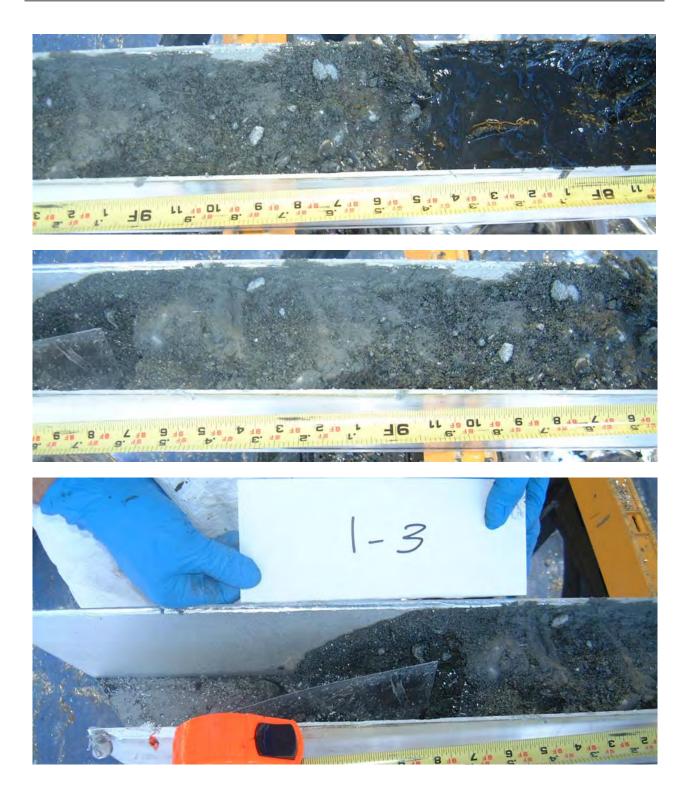
Photographs of NAPL Investigation Core 1-2: Starting at the top of the core



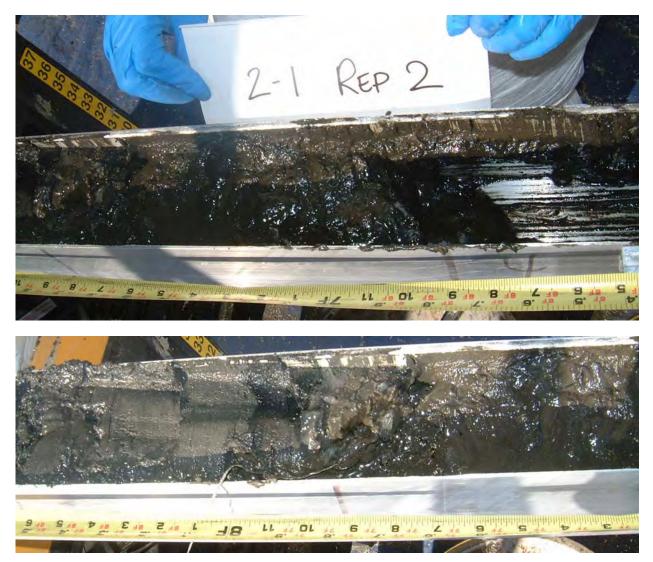


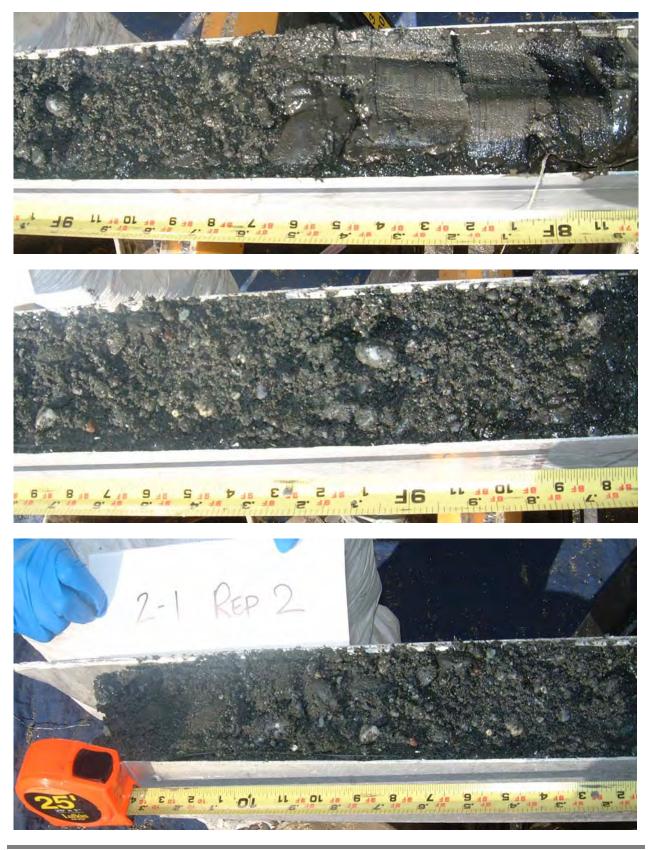
Photographs of NAPL Investigation Core 1-3: Starting at the top of the core



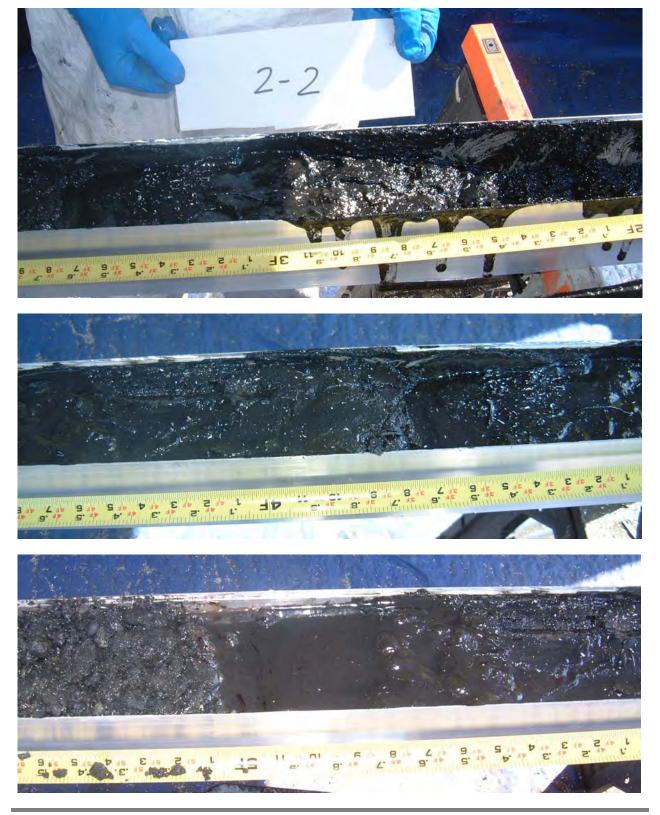


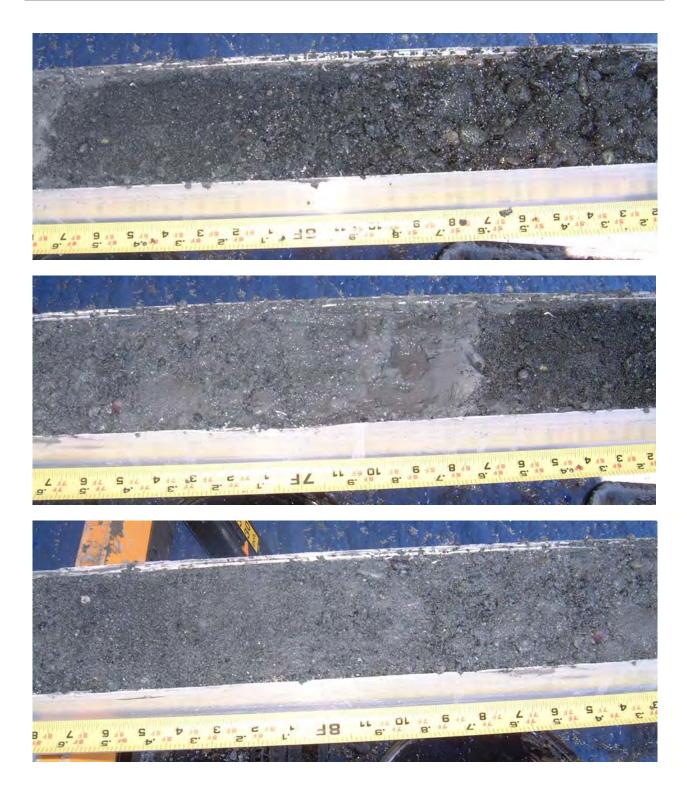
Photographs of NAPL Investigation Core 2-1 Rep 2: Starting at the top of the core



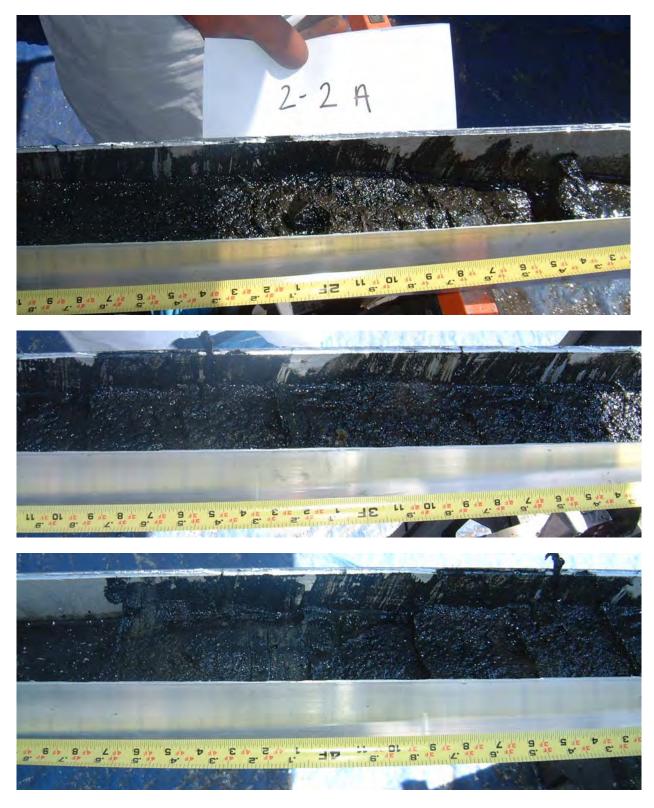




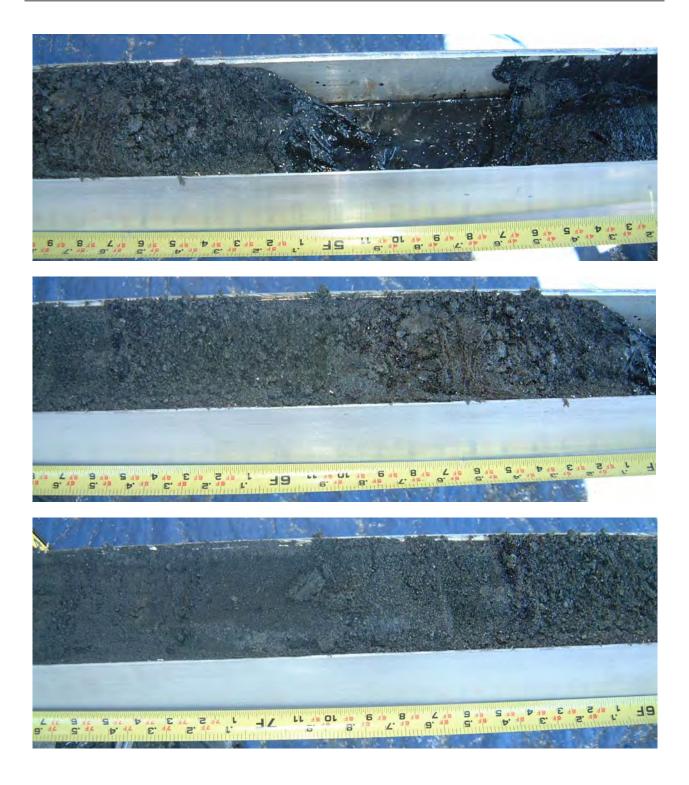


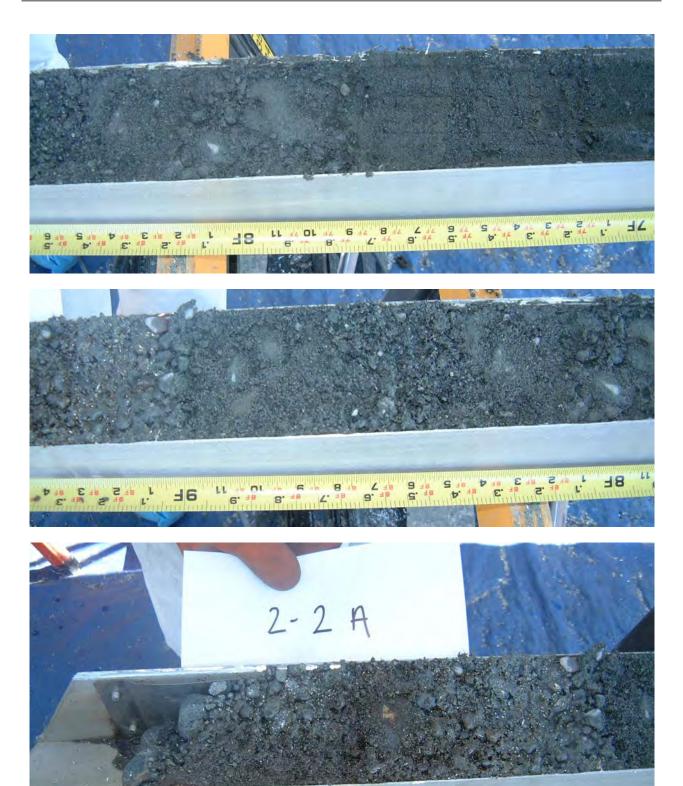






Photographs of NAPL Investigation Core 2-2 A: Starting at the top of the core





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OL

2

LL

6

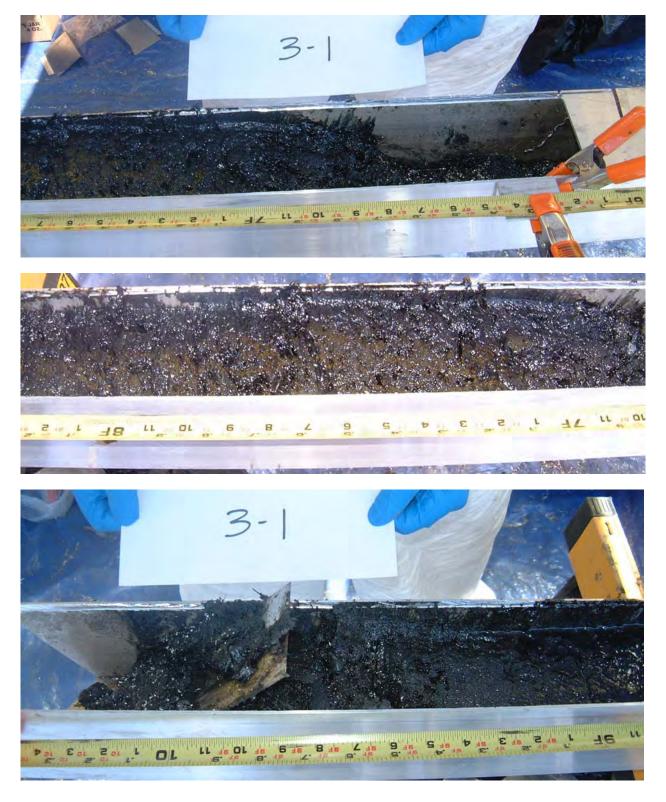
OL

8

36

LL

Photographs of NAPL Investigation Core 3-1: Starting at the top of the core





Photographs of NAPL Investigation Core 3-2: Starting at the top of the core



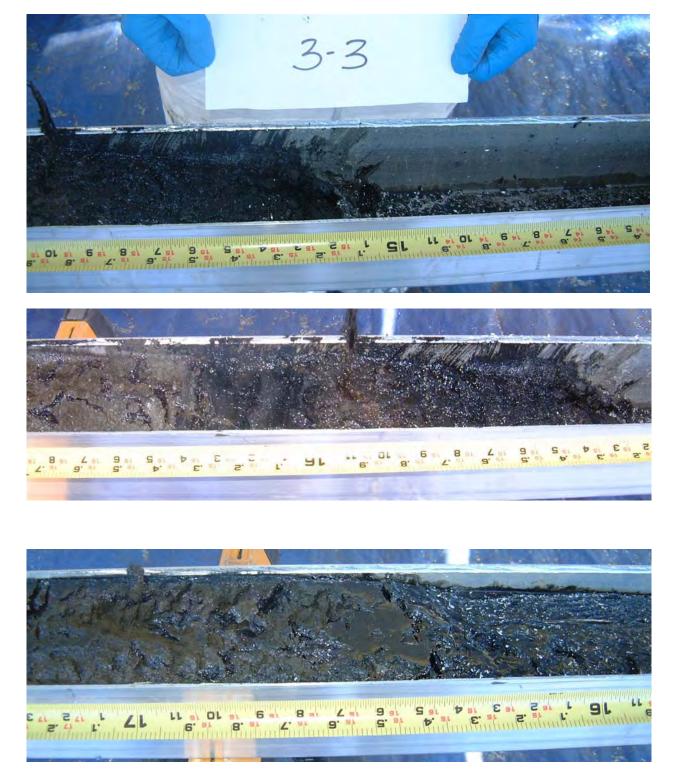


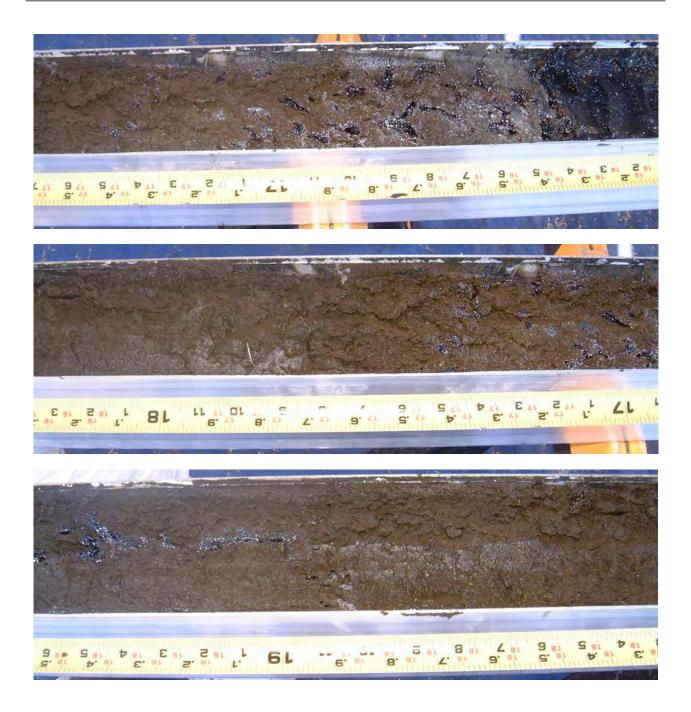


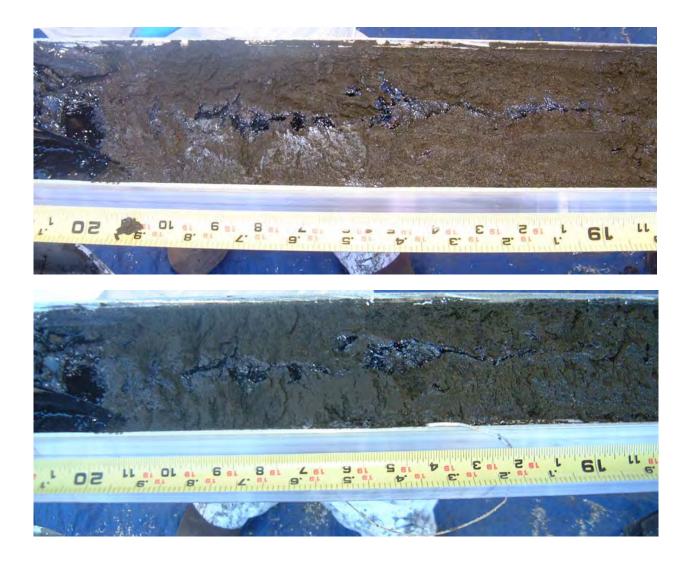






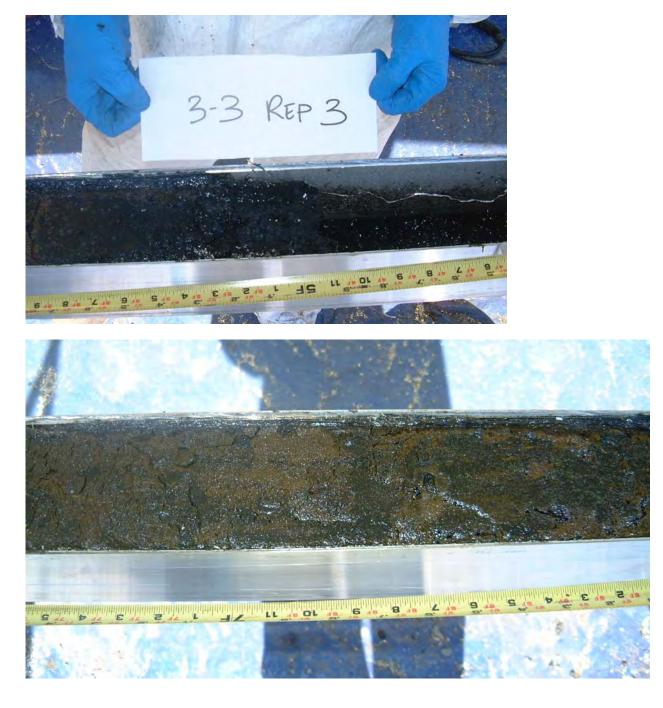




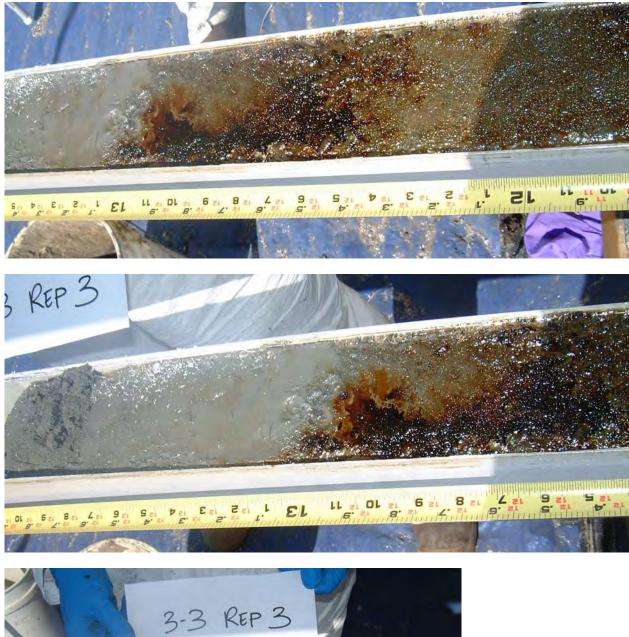




Photographs of Core 3-3 Rep 3: Starting at the top of the core









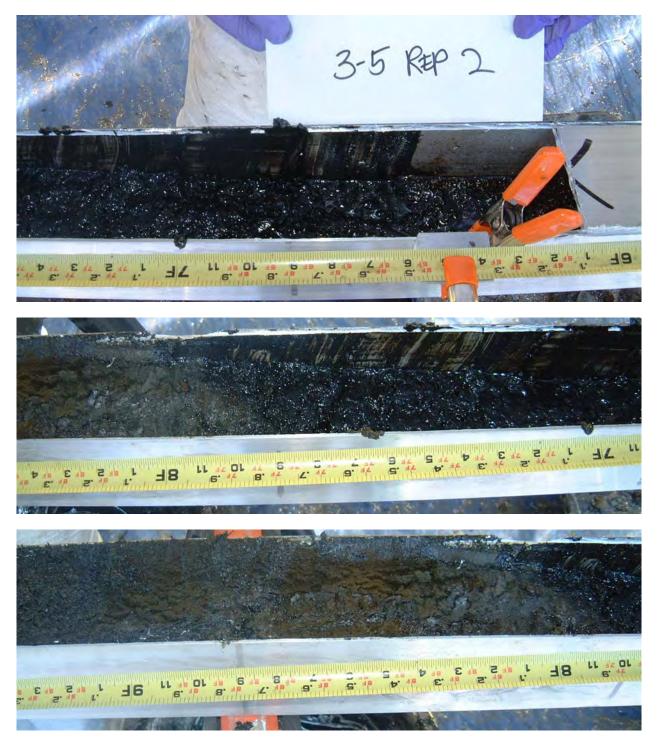
Photographs of NAPL Investigation Core 3-4: Starting at the top of the core







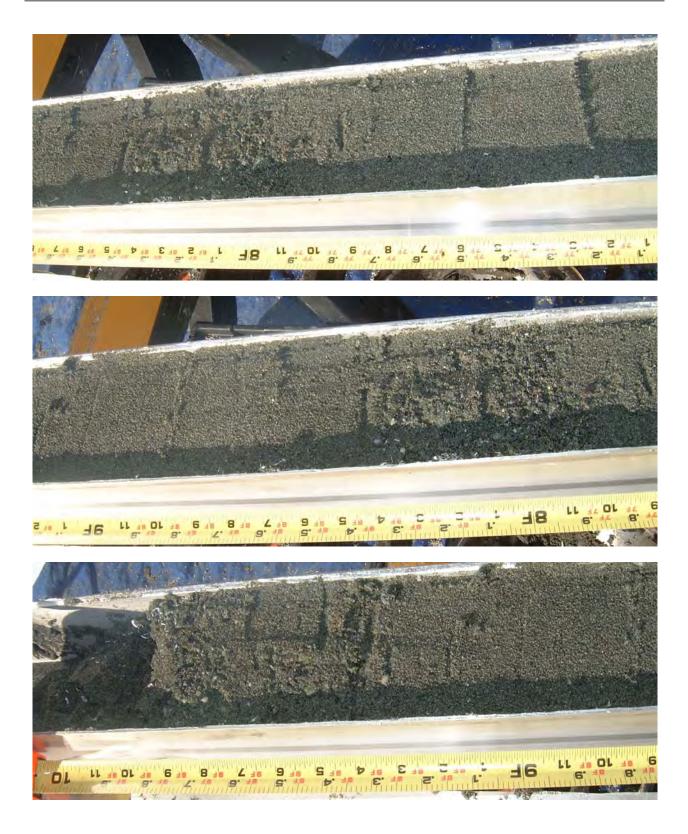
Photographs of NAPL Investigation Core 3-5 Rep 2: Starting at the top of the core

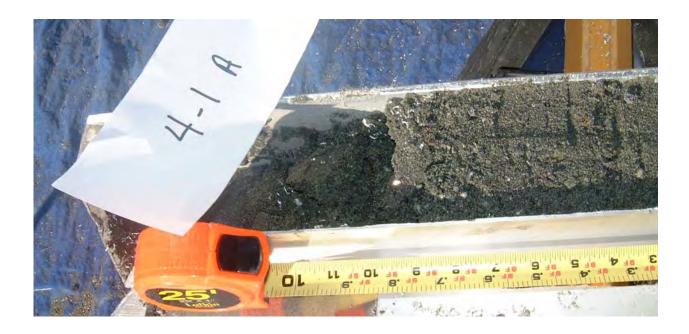




Photographs of NAPL Investigation Core 4-1 A: Starting at the top of the core



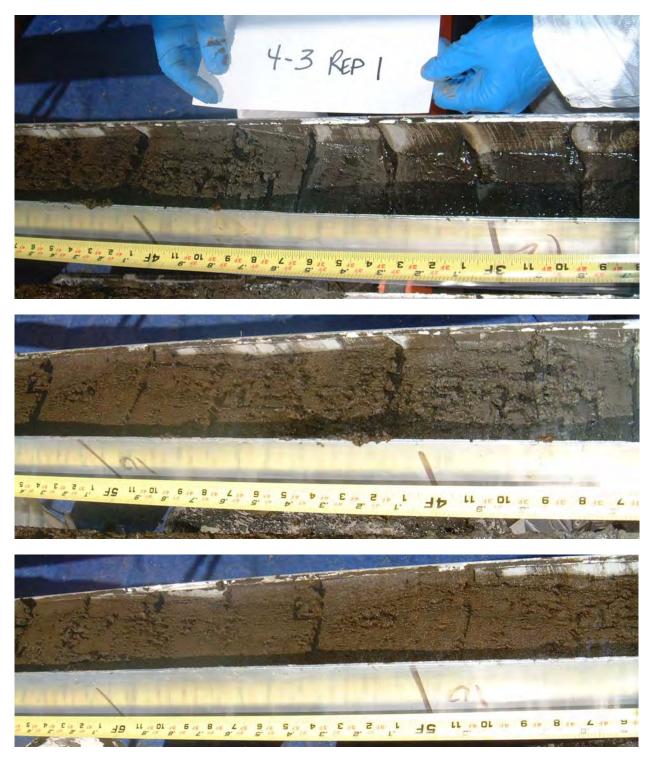


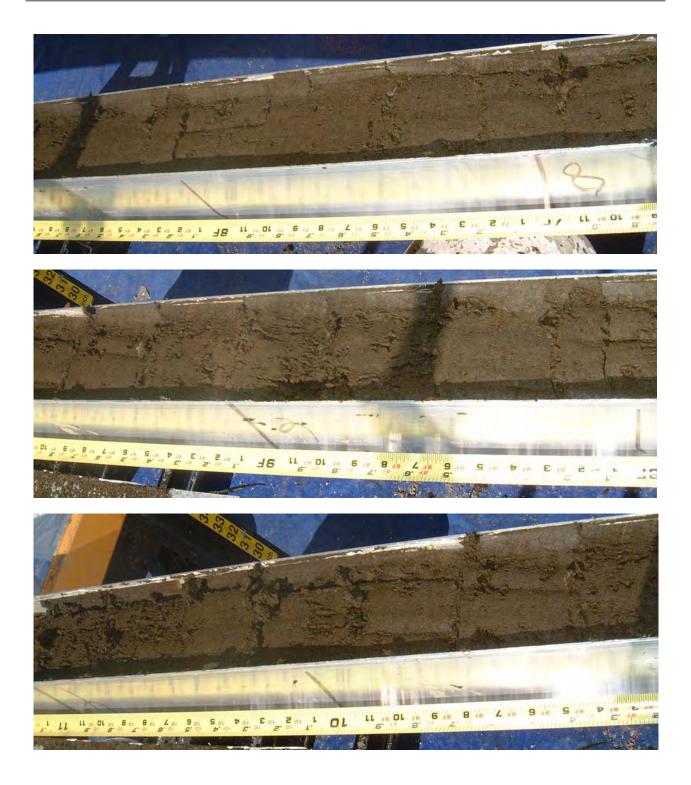


Photographs of NAPL Investigation Core 4-2: Starting at the top of the core



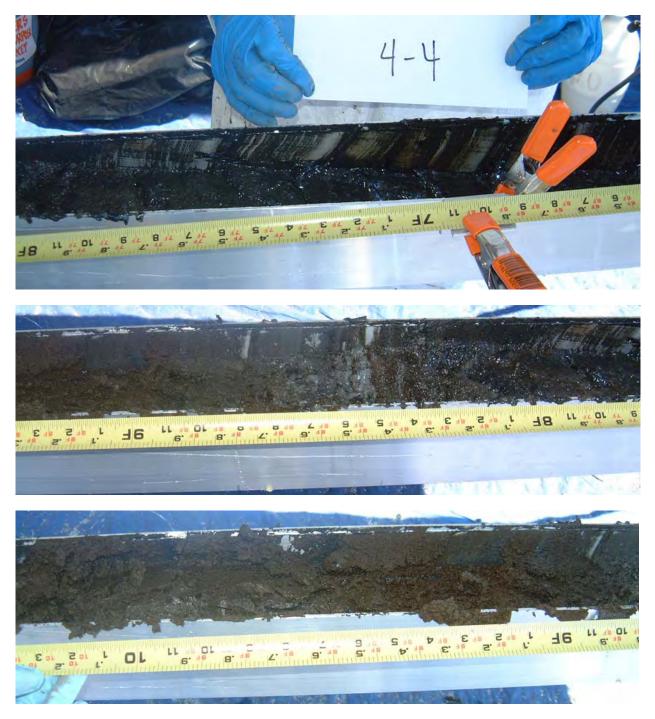
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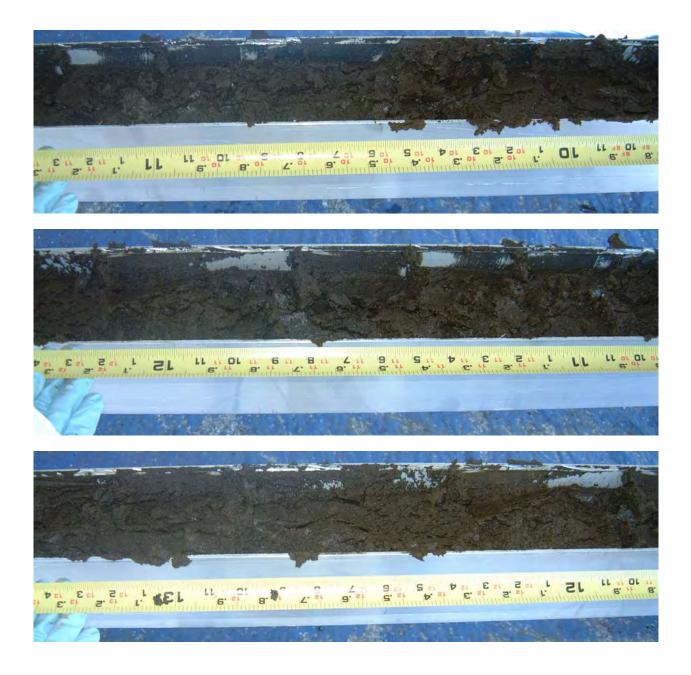


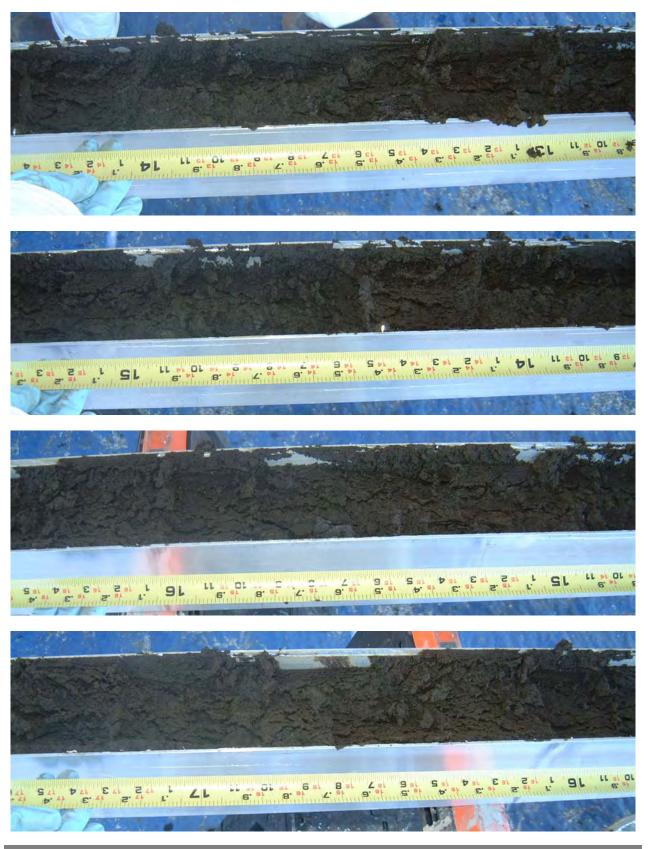


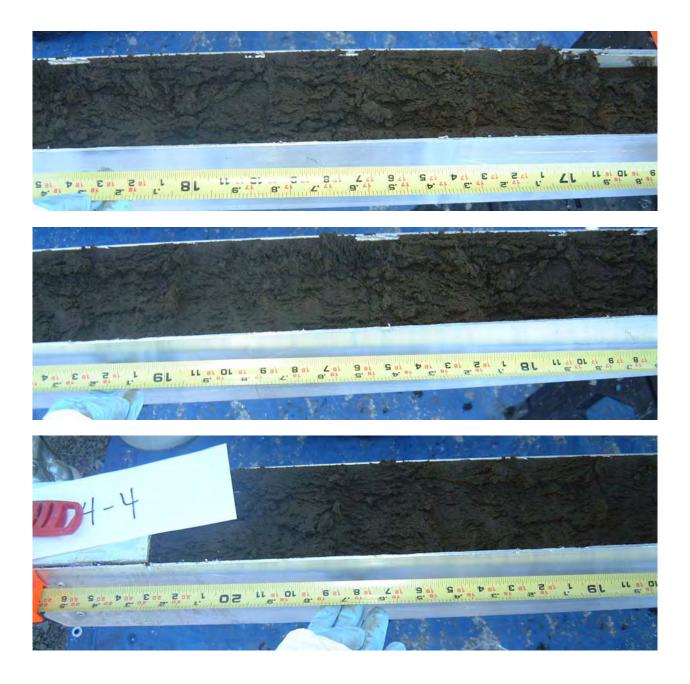


Photographs of NAPL Investigation Core 4-4: Starting at the top of the core

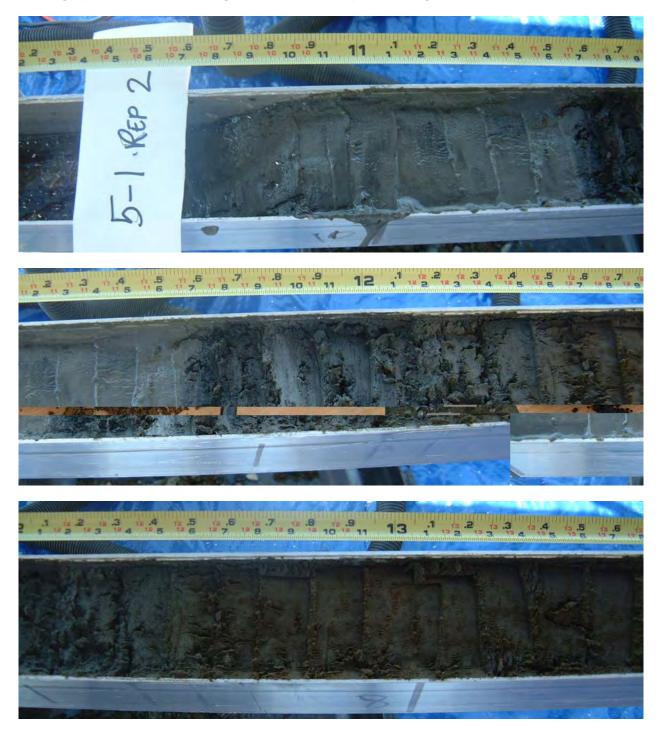








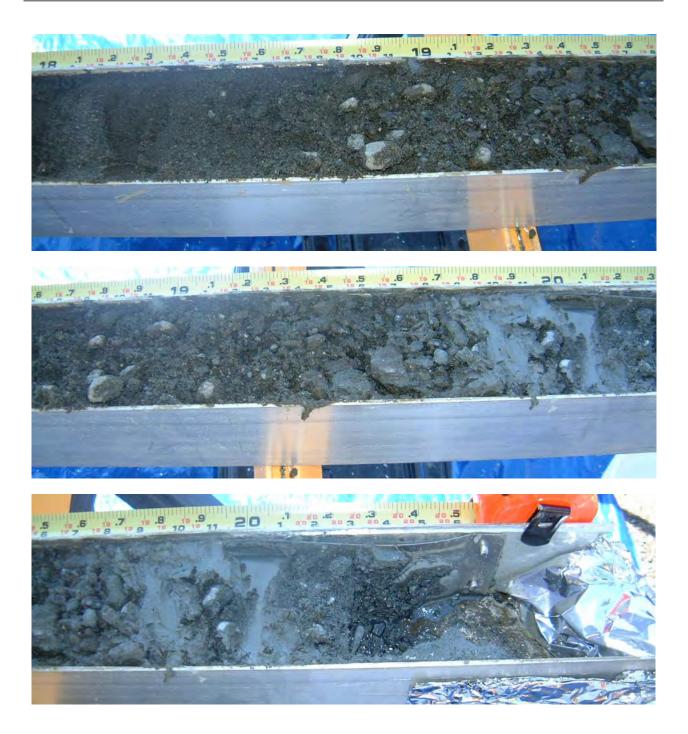
Photographs of NAPL Investigation Core 5-1 Rep 2: Starting at the top of the core



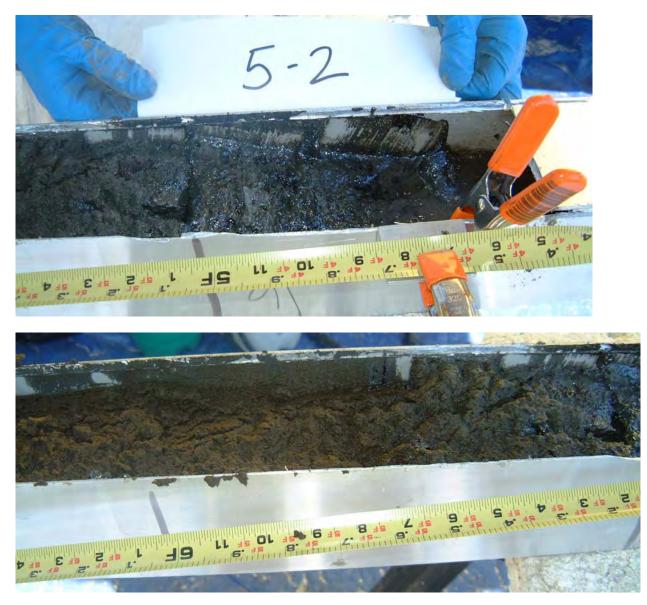








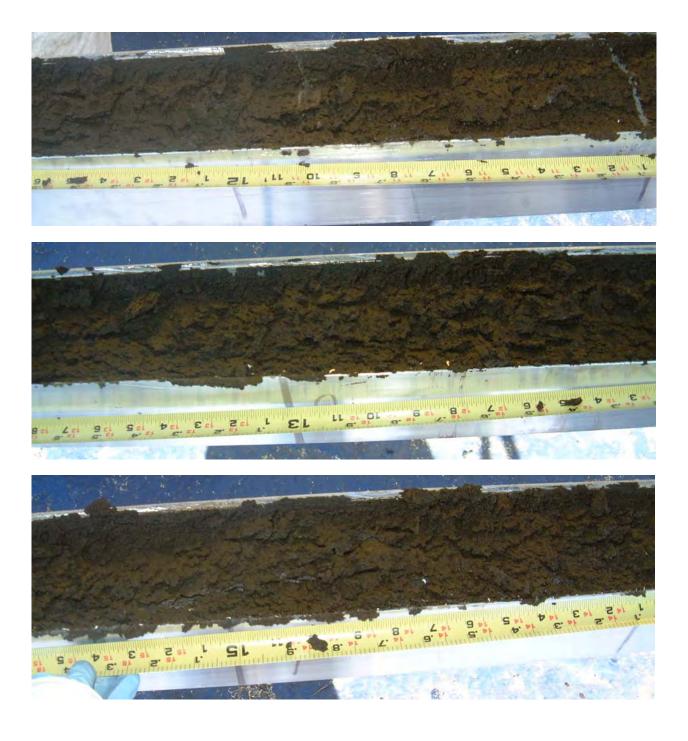
Photographs of NAPL Investigation Core 5-2: Starting at the top of the core





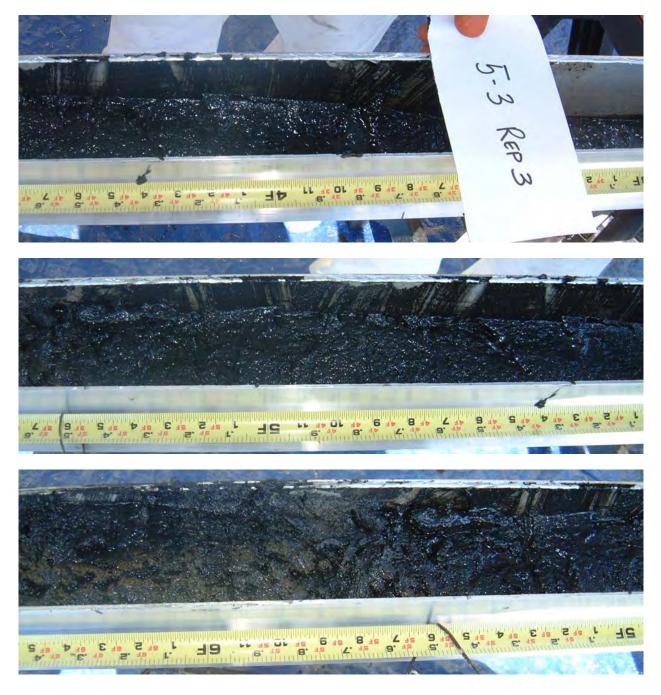


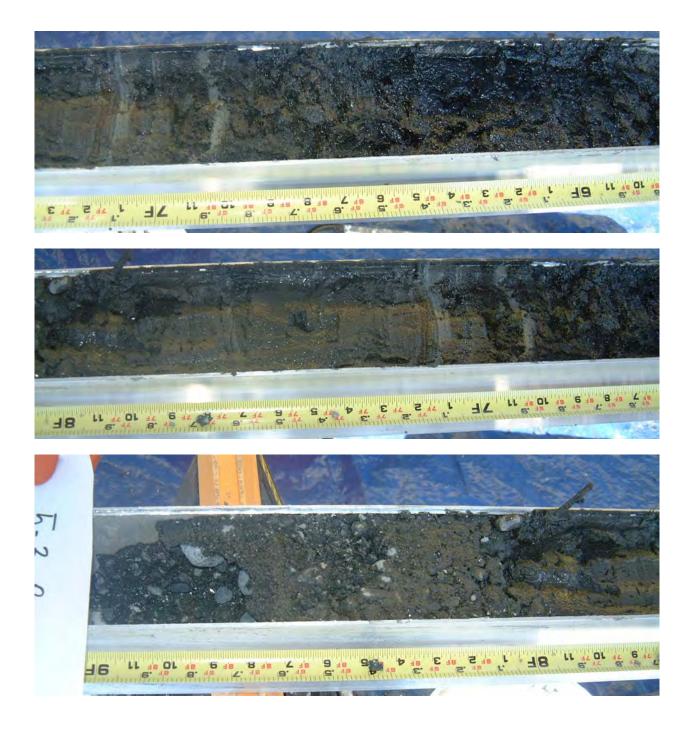






Photographs of NAPL Investigation Core 5-3 Rep 3: Starting at the top of the core

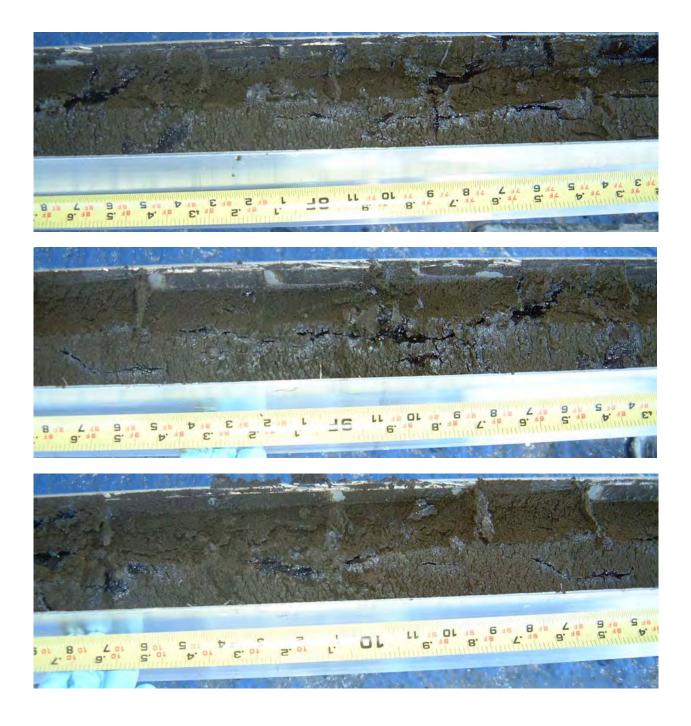


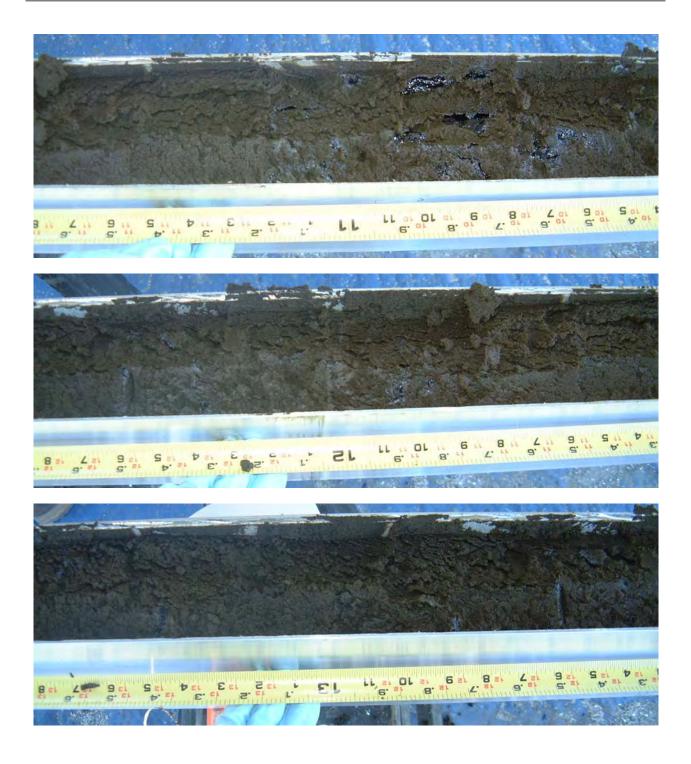




Photographs of NAPL Investigation Core 5-4: Starting at the top of the core









ATTACHMENT 2D-5 Battelle February 2007 Data Report

Battelle The Business of Innovation

Data Report

Gas Works Park Soils February 2007

Prepared for:

Brad Bessinger Exponent, Incorporated

Prepared by:

Battelle 397 Washington Street Duxbury, Massachusetts 02332 Chain of Custody

ShpNo SHP-070116-01

The Business of	Business of Innovation Battelle Project No: 06746-0001										
Sample Receip	ot Form										
					Approve	d: 📃 Auth	orized 📃				
Project Number:	BN03005.001		Client:	Exponent							
Received by:	Seyfert, Jeannine	9	Date/Ti	me Received:	Tuesday, January 16, 20	007 10:44 AM					
No. of Shipping Con	tainers: 1	_									
SHIPMENT											
Method of Delivery:	Commercial Car	rier	Tracki	ing Number:	8585-3984-8924						
COC Forms:	Shipped w	ith samples	No For	ms							
Cooler(s)/Box	x(es)										
Cntr Type	Tracki	ng No.	Seal	Seal Condition			Smps				
1 of 1 Cooler			Таре	Intact	Intact	3.0	22				
Samples											
Sample Labels:		Sample label	s agree with	n COC forms							
		Discrepancie	s (see Samj	ple Custody Corr	rective Action Form)						
Container Seals:		Tape C	ustody Sea	ls 🗌 Other Se	als (See sample Log)						
		Seals intact for	or each ship	pping container							
		Seals broken	(See sampl	e log for impact	ed samples)						
Condition of Samples	:	Sample conta	iners intac	ŀ							
· · · · · · · · · · · · · · · · · · ·					Custody Corrective Actio	n Form)					
					·						
Temperature upon re	-				Yes No						
(Note: If temperature u	ipon receipt diffe	rs from required	conditions,	see sample log o	comment field)						
Samples Acidified:		Yes No	✔ Unk	nown							
Initial pH 5-9?:		Yes No	🖌 NA								
If no, individual sampl	e adjustments on	the Auxiliary San	nple Receip	ot Form							
Total Residual Chlori	ing Procont?.	Yes No	✓ NA								
If yes, individual samp				pt Form							
Head Space <1% in s	-	-	Yes Yes		NA						
Individual sample devi	ations noted on s	ample log									
Samples Containers:											
Samples returned in PC	C-grade jars:	Yes No	🖌 Unk	nown /Lot No.:	Unknown						
Storage Location:	Field Roor	n II: Refrigerator	- R0004 (Lo	ower Col BDO	IDs Assigned: R55	21 - R5542					
Samples logged in by	y: Seyfert, Je	eannine			Date/Time:	01/16/2007 10):44 AM				
Approved By:					Approved On:						
Authorized By:					Authorized On:		-				

ShpNo SHP-070116-01

Battelle Project No: 06746-0001

Sample Receipt Form Details

Client: Exponent

BN03005.001

Project Number:

Approved: Authonized

Received by:	ed by: Seyfert, Jeannine		ate/Time	Received: Tue	Date/Time Received: Tuesday, January 16, 2007 10:44 AM	07 10:44 AN	~		1			
No. of :	No. of Shipping Containers:	-										
BDO Id:	BDO Id: Client Sample ID:	Collection Date:	ate:	Login Date:	Ctrs: Matrix:	Temp	Temp: pH: TRC: VOC:	TRC	007	: Stored In:	Loc:	No: Comments:
R5521	GWP07T01	01/11/07 12:00	00:	01/16/07 11:04	1 TAR	с	ΝA	ΝA	ΝA	R0004 (Lower C	BIN	21
R5522	GWP07DW401	01/11/07 10:54	:54	01/16/07 11:05	1 NAPL	с С	ΝA	ΝA	ΝA	R0004 (Lower C	BIN	21
R5523	GWP07DW402	01/11/07 10:54	:54	01/16/07 11:06	1 NAPL	с	ΝA	ΝA	NA	R0004 (Lower C	BIN	21
R5524	GWP07MW9	01/12/07 10:50	:50	01/16/07 11:07	1 NAPL	с	٩N	٩N	AA	R0004 (Lower C	BIN	21
R5525	GWP07T02	01/11/07 12:10	:10	01/16/07 11:09	1 TAR	ო	ΝA	ΝA	NA	R0004 (Lower C	BIN	21
R5526	GWP07T03	01/11/07 12:10	10	01/16/07 11:19	1 TAR	ო	ΝA	ΝA	NA	R0004 (Lower C	BIN	21
R5527	GWP07T04	01/11/07 12:20	:20	01/16/07 11:20	1 TAR	ო	ΝA	ΝA	NA	R0004 (Lower C	BIN	21
R5528	GWP07T05	01/11/07 12:35	:35	01/16/07 11:21	1 TAR	ო	ΝA	ΝA	NA	R0004 (Lower C	BIN	21
R5529	GWP07T06	01/11/07 12:40	:40	01/16/07 11:22	1 TAR	с	٨A	٨A	AN	R0004 (Lower C	BIN	21
R5530	GWP07T07	01/11/07 12:45	:45	01/16/07 11:23	1 TAR	с	٩N	٩N	AN	R0004 (Lower C	BIN	21
R5531	GWP07T08	01/11/07 13:10	:10	01/16/07 11:28	1 TAR	с	ΝA	ΝA	NA	R0004 (Lower C	BIN	21
R5532	GWP07T09	01/11/07 13:40	:40	01/16/07 11:28	1 TAR	ო	ΝA	ΝA	NA	R0004 (Lower C	BIN	21
R5533	GWP07T10	01/11/07 13:48	:48	01/16/07 11:29	1 TAR	ო	ΝA	ΝA	NA	R0004 (Lower C	BIN	21
R5534	GWP07T11	01/11/07 14:20	:20	01/16/07 11:31	1 TAR	ო	ΝA	ΝA	NA	R0004 (Lower C	BIN	21
R5535	GWP07T12	01/11/07 14:50	:50	01/16/07 11:33	1 TAR	ო	٨A	ΝA	NA	R0004 (Lower C	BIN	21
R5536	GWP07T13	01/11/07 14:55	:55	01/16/07 11:33	1 TAR	e	٨A	ΝA	NA	R0004 (Lower C	BIN	21
R5537	GWP07T14	01/11/07 15:05	:05	01/16/07 11:34	1 TAR	ო	٨A	ΝA	NA	R0004 (Lower C	BIN	21
R5538	GWP07S01	01/12/07 9:30	30	01/16/07 11:35	1 TAR	ю	٨A	ΝA	NA	R0004 (Lower C	BIN	21
R5539	GWP07S02	01/12/07 9:35	35	01/16/07 11:36	1 TAR	e	٨A	ΝA	NA	R0004 (Lower C	BIN	21
R5540	GWP07S03	01/12/07 10:15	:15	01/16/07 11:36	1 TAR	e	٨A	ΝA	ΝA	R0004 (Lower C	BIN	21
R5541	GWP07S04	01/12/07 10:30	:30	01/16/07 11:37	1 TAR	e	٨A	ΝA	ΝA	R0004 (Lower C	BIN	21
R5542	TDW3-4.5	09/26/06 12:56	:56	01/16/07 11:39	1 SOIL	3	NA	NA	NA	R0004 (Lower C	BIN	21
Total S	Total Samples: 22											

Page of 2	E ^x ponent		ן פוגס	, enta	uuoui,	rchi/		85521	R5522	R5523	RSS24	RSSZS	R5526	R5527	R5528	R5529	R5530	R5531	R5532	R5533	R5534	R5535	R5536	R5537	R5533	R5539	R5540			Custody Seal Intact: Ves No None	Date/Time: 11307 13:32	all that and the second second second	
_YSIS REQUEST FORM	2005,00	B. Bessinger 1. Massingale (Floyd Snider)	ses Requested	per bel	5-2 	9088 V R	EFP	×	X	X	X	X	X	X	X	×	×	X	×	X	×	X	×	×	×	×				Condition of Samples Upon Receipt:	21	101 10	
CHAIN OF CUSTODY RECORD/SAMPLE ANAL	Project: (Name and Number) GAS WORKS PARK, BN03	Office: BE Samplers:	PPLIED	HINGTON ST.	_	V de taca	Sample No. Tag No. Date Time Matrix O	GWP07T01 75601 111/0712:00 T X	GWP07DW401 75650 1/1/07 10:54 NAPL X	(auport Du4021 75 65 1/1/07 10:54 NAPL X	75625/12107 10:50	GWP07T02 75602 1/1/0712:10 T X	X T 01:21/07/17 75603 1/11/07 12:10 T X	1	("WP07TD5 75605 /11/12:35 T X			TO8 75608 1/1/67	170/11/1 POJZC POTT	7710 756 10 1/1/071	7711 75611 1/1/07	1712 75612 1/11/07	6WP 07T13 75613 1/1/07 14:55 T X) ~	6WP07501 75621 1/12/07 04:30 T X	6WP07502 75622 1/12/07 04:35 T X	GWP07503 75623 1/12/07 10:15 T X	Matrix GW - Groundwater SL - Soil SD - Sediment SW - Surface water Code:	OTHER - Please identify codes T-TAR	Shipped FedFx/LPS Courier Other	These Arne (Spann	Heinquished by: DAU ANYTYU ANY (Signature)	

	05, 00	B. Bessinger / J. Mussingle (Floyd Snider) 3	פוגי		Autor Samo Samo Mon Mon Mon Mon Mon Mon Mon Mon Mon Mo	-5 d -0121	ES ES ES ES EX Extern	X X X 85541	X X A A A A A A A A A A A A A A A A A A							water Priority: Normal Bush time period	Condition of Samples Custody Seal Intact: Ves No None	2	ime: 1/15/07 1330 Received by: Jeannin Series deyland. Date/Time: 1/16/07 10:40	
APLE ANALYSIS REQUEST FORM	BN050	Samplers: R. Bessivier /	Analyses Rec	O N V	0413 -202	W W W	Eb 29	T X X								Normal	Condition of Samples Upon Receipt:	13:32 Received by:	707 1330 Received by:	1
<u>U</u>	(Name and Number) CAS WORKS PARK,	Exponent Contact: BRAD BESSINGER Office: BE	Ship to: BATTELLE APPLIED ENVIRONMENTAL CHEMILITY	397 WASHINGTON STREET		NY	Time	6WP07504 75624 Y12/07 10:33 7	GTDW3-45 - 91,2610K 1254 Soul							Matrix GW - Groundwater SL - Soil SD - Sediment SW Odde: OTHER - Please identify codes T-TAR	Shipped FedEx/UPS Courier Other	ed by: Kal Kenne	lessilla	(Signature)

SHC Data and Chromatograms

SHC and TPH – Sediment QA/QC Summary Batches 07-0010

PROJECT:	Exponent – Gas Works Park
PARAMETER:	Saturated Hydrocarbons (SHC) and Total Petroleum Hydrocarbons (TPH)
LABORATORY:	Battelle, Duxbury, MA
MATRIX:	Sediment
SAMPLE CUSTODY:	Eighteen tar samples, three NAPLs samples, and 1 soil sample were received at the Battelle
	Duxbury Operations (BDO) Laboratory on 1/16/2007. Upon receipt of samples, the
	temperatures of the coolers were taken and the samples were logged into the laboratory and
	given unique IDs. The temperature of the cooler upon receipt was within the acceptable
	range. Samples were either stored in an access-limited walk-in refrigerator at 4°C until
	sample preparation could begin. The soil and tar samples were extracted together in one
	analytical batch, batch 07-0001.

QA/QC DATA QUALITY OBJECTIVES:

	Reference Method	Blank	Surrogate Recovery	LCS/MS Recovery	MS/MSD Precision	Control Oil % Diff.
SHC and TPH	General NS&T	< 5x MDL	40-120% Recovery	40-120% Recovery MS target spike must be > 5 x background	< 30% RPD	PD < 30% for 90% of the analytes

METHOD:

Soil and tar samples were extracted following general NS&T methodologies.

Approximately 5-8 grams of sample was spiked with SHC, PAH, and biomarker surrogates and serially extracted three times with dichloromethane using orbital shaker table techniques. The combined extracts were dried over sodium sulfate and concentrated by Kuderna-Danish and nitrogen evaporation techniques. The sample extracts were split in half; one-half of the extract was removed for archiving; the other half was processed through an alumina gravity column to isolate the hydrocarbon fractions of interest. The weights of the resulting extracts were determined gravimetrically. The extracts were concentrated to 1 ml, split, and spiked with IS. The pre-injection volume and/or extract split were adjusted to achieve 5 mg/mL. One extract was submitted for PAH and petroleum biomarker analysis, and the second extract was submitted for SHC and TPH analysis.

SHC and TPH were measured by gas chromatography with flame ionization detection (GC/FID). An initial calibration consisting of target analytes was completed prior to analysis to demonstrate the linear range of analysis. Calibration verification was performed at the beginning and end of each 12 hour period in which samples were analyzed. Concentrations of SHC and TPH were calculated by the internal standard method. Normal alkanes were quantified using the average RF generated from the initial calibration. TPH concentrations were quantified using the average RF of nC8 through nC40. Isoprenoid hydrocarbon concentrations were quantified using the average RF of nC8 through nC40. Isoprenoid hydrocarbon and immediately following each target isoprenoid hydrocarbon.

HOLDING TIMES: Samples were prepared for analysis in one analytical batch. Samples were extracted within 14 days of sample receipt and analyzed within 40 days of extraction. Holding times from collection to extraction for soil samples are 14-days if refrigerated, 365-days if frozen. Since the storage conditions for sample TDW3-4.5, collected 9/26/2006, prior to its arrival at BDO are unknown, this data has been conservatively qualified with a "T".

Batch ID	Extraction Date	Analysis Date(s)
07-0010	1/24/2007	1/31/2007 - 2/1/2004

SHC and TPH – Sediment QA/QC Summary Batches 07-0010

PROCEDURAL BLANK (PB):	A procedural blank (PB) was prepared with each analytical batch. The blank was analyzed ensure the sample extraction and analysis methods were free of contamination.							
	07-0010 – No exceedences noted.							
	Comments- None							
LABORATORY CONTROL SAMPLE	A laboratory control sample (LCS) was prepared each analytical batch. The percent recoveries of target analytes were calculated to measure data quality in terms of accuracy.							
(LCS):	07-0010 – No exceedences noted.							
	Comments – None.							
SURROGATE RECOVERY:	Two surrogate compounds were added prior to extraction, including o-terphenyl and 5a- androstane. The recovery of the surrogate compound was calculated to measure data quality in terms of accuracy (extraction efficiency).							
	07-0010 – No exceedences noted.							
	Comments – None							
CONTROL OIL:	A control oil (North Slope Crude) was prepared with the analytical batch. The percent difference (PD) between the measured value and the target value was calculated to measure data quality in terms of accuracy.							
	07-0010 – No exceedences noted.							
	Comments – None.							
CALIBRATIONS:	The GC/FID is calibrated with a minimum 5 level curve for all compounds. The percent relative standard deviation (%RSD) between RF for the individual target analytes must be $\leq 25\%$, and the mean RSD of all target analytes must be $< 20\%$. Each batch of samples analyzed is bracketed by continuing calibration verification (CCV) sample, run at a frequency of minimally every 12 hours. The PD between the true value and the CCV should be $< 20\%$ for individual analytes. Additionally an initial calibration check (ICC) sample is run immediately after each initial calibration. The PD between the ICC and the initial calibration should be $< 20\%$.							
	07-0010 – No exceedences noted.							

Comments – None.

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07T01	GWP07T02	GWP07T03	GWP07T04
Battelle ID	R5521-P	R5525-P	R5526-P	R5527-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/11/07	01/11/07
Extraction Date	01/24/07	01/24/07	01/24/07	
				01/24/07
Analysis Date	01/31/07	01/31/07	01/31/07	02/01/07
Analytical Instrument	FID	FID	FID	FID
% Moisture	28.88	50.1	50.3	34.93
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.44	1.02	1.01	1.44
Size Unit-Basis	G_DRY	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit	55.58	245.10	198.1	92.67
Units	MG/KG_DRY	MG/KG_DRY	MG/KG_DRY	MG/KG_DRY
n-Nonane	U	U	U	U
n-Decane	U	1193.55	996.66	444.11
n-Undecane	U	612.43	533.37	88.93 J
	948.54			
n-Dodecane		2935.42	2429.74	1312.88
n-Tridecane	468.84 ME	2504.47 ME	2367.65 ME	1524.69 ME
Isoprenoid RRT 1380	U	U	U	U
n-Tetradecane	152.15	336.6	264.56	193.36
Isoprenoid RRT 1470	136.99	278.2	231.91	176.53
n-Pentadecane	79.8	159.6 J	131.08 J	98.43
n-Hexadecane	108.22	258.18	177.19 J	125.75
Norpristane (1650)	201.38	404.98	319.4	240.35
n-Heptadecane	52.84 J	134.02 J	95.89 J	65.46 J
Pristane	55 J	121.46 J	86.69 J	62.31 J
n-Octadecane	1472.34	3144.3	2446.92	1737.93
Phytane	36.74 J	76.6 J	60.75 J	37.48 J
n-Nonadecane	11.78 J	29.82 J	21.48 J	16.34 J
n-Eicosane	37.55 J	82.28 J	72.9 J	43.75 J
n-Heneicosane	454.04	1059.47	826.47	581.65
n-Docosane	157.29	340.27	233.82	196.96
n-Tricosane	38.99 J	82.85 J	63.45 J	43.91 J
n-Tetracosane	80.7	198.52 J	138.97 J	91.93 J
n-Pentacosane	U	U	528.12	U
n-Hexacosane	U	U	176.8 J	U
n-Heptacosane	U	U	U	U
n-Octacosane	U	U	U	U
n-Nonacosane	U	U	U	U
n-Triacontane	U	U	U	U
n-Hentriacontane	U	U	U	U
n-Dotriacontane	U	U	U	U
n-Tritriacontane	U	U	U	U
n-Tetratriacontane	U	U	U	U
n-Pentatriacontane	U	U	U	U
n-Hexatriacontane	U	Ŭ	Ŭ	U
n-Heptatriacontane	Ŭ	Ŭ	Ŭ	Ŭ
n-Octatriacontane	Ŭ	Ū	Ŭ	Ŭ
n-Nonatriacontane	U	U	U	U
n-Tetracontane	Ŭ	Ŭ	Ŭ	U
TPH(total)	194301.96	600370.63	457694.31	232050.17
	10-1001.00	000070.00	-01007.01	202000.17
Surrogate Recoveries (%)				
O-Terphenyl	91	106	99	108
5a-androstane	84	90	87	88

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07T05	GWP07T06	GWP07T07	GWP07T08
Battelle ID	R5528-P	R5529-P	R5530-P	R5531-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/11/07	01/11/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/01/07	02/01/07	02/01/07	02/01/07
Analytical Instrument	FID	FID	FID	FID
% Moisture	4.23	6.94	6.47	30.28
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.92	1.93	1.87	1.49
Size Unit-Basis	G_DRY	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit	270.83	174.16	178.3	167.92
Units	MG/KG_DRY	MG/KG_DRY	MG/KG_DRY	MG/KG_DRY
n-Nonane	U	U	U	U
n-Decane	U	Ŭ	Ŭ	U
n-Undecane	Ŭ	Ŭ	Ŭ	Ŭ
n-Dodecane	154.79 J	102.49 J	132.16 J	750.35
n-Tridecane	958.19 ME	709.86 ME	887.45 ME	2733.76 ME
Isoprenoid RRT 1380	U	U U	U	2700.70 ML
n-Tetradecane	349.29	280.5	314.44	290.87
Isoprenoid RRT 1470	224.99 J	194.69	205.67	572.59
n-Pentadecane	75.85 J	57.61 J	55.65 J	236.74
n-Hexadecane	184.63 J	165.06 J	177.34 J	63.43 J
Norpristane (1650)	673.76	641.75	640.32	48.89 J
n-Heptadecane	142.89 J	104.99 J	129.55 J	48.89 J 140.64 J
Pristane	124.44 J	117.49 J	135.72 J	223.04
n-Octadecane	7021.65	6582	6697.6	3557.74
Phytane	847.82	798.53	885.03	3557.74 38.47 J
n-Nonadecane	047.02 U	790.55 U	005.05 U	30.47 J U
n-Eicosane	246.16 J	228.46	206.47	U
n-Heneicosane	246.16 J U	220.40 U	200.47 U	1108.3
n-Docosane	876.93	767.75	896.07	
n-Tricosane	183.14 J	U	090.07 U	639.18 U
n-Tetracosane	393.55	295.92	U	527.55
n-Pentacosane	1499.75	174.84	1516.28	527.55 U
n-Hexacosane	1156.24	1079.68	1269.93	U
	U	U 1079.00	1209.95 U	U
n-Heptacosane n-Octacosane	4408.6	3893.14	4634.83	U
n-Nonacosane	4408.8 U	3693.14 U	4034.03 U	U
n-Triacontane	228.82 J	178.94	U	U
n-Hentriacontane	220.02 J U	U	U	U
n-Dotriacontane	U	U	U	U
n-Tritriacontane	260.88 J	207.53	U	U
n-Tetratriacontane	200.88 J 214.32 J	207.55	U	U
n-Pentatriacontane		220.89 U	U	U
	U		U	U
n-Hexatriacontane n-Heptatriacontane	U U	U U	U	U
	U	U	U	U
n-Octatriacontane				-
n-Nonatriacontane	U	U	U	U
n-Tetracontane	U	U	U	U
TPH(total)	297041.61	269380.89	292690.39	227878.18
Surrogate Recoveries (%)				
O-Terphenyl	101	117	102	97
5a-androstane	108	111	103	81

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07T09	GWP07T10	GWP07T11	GWP07T12
Battelle ID	R5532-P	R5533-P	R5534-P	R5535-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/11/07	01/11/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/01/07	02/01/07	02/01/07	02/01/07
Analytical Instrument	FID	FID	FID	FID
% Moisture	19.46	32.51	5.51	10.85
		NA	NA	NA
% Lipid Matrix	NA TAR	TAR	TAR	TAR
Matrix Sample Size	1.72	1.45	2.05	1.79
•	G_DRY	G_DRY	G_DRY	G_DRY
Size Unit-Basis	193.8	137.93	19.51	74.68
Minimum Reporting Limit Units	MG/KG_DRY	MG/KG_DRY	MG/KG_DRY	MG/KG_DRY
Onits	MG/RG_DRT	MG/KG_DKT	MG/KG_DR1	
n-Nonane	U	U	U	U
n-Decane	Ŭ	Ŭ	3.13 J	70.47 J
n-Undecane	Ŭ	505.29	6.18 J	50.98 J
n-Dodecane	564.36	1619.57	5.1 J	195.2
n-Tridecane	4528.19 ME	6168.86 ME	154.92 ME	1161.6 ME
Isoprenoid RRT 1380	U	U	U	U
n-Tetradecane	296.14	245.96	94.21	104.69
Isoprenoid RRT 1470	604.8	223.3	65.08	143.44
n-Pentadecane	258.95	119.69 J	20.29	70.38 J
n-Hexadecane	386.22	172.46	42.1	122.55
Norpristane (1650)	59.23 J	301.71	25.35	344.25
n-Heptadecane	234.55	86.53 J	27.6	59.62 J
Pristane	319.15	105.86 J	49.55	69.94 J
n-Octadecane	3992.53	2285.79	350.75	2343.65
Phytane	56.2 J	54.28 J	10.08 J	84.67
n-Nonadecane	U	U	U	U
n-Eicosane	Ŭ	73.73 J	Ŭ	79.21
n-Heneicosane	1428.89	743.82	160.46	806.59
n-Docosane	782.6	274.01	84.28	329.85
n-Tricosane	U	U	U	60.23 J
n-Tetracosane	670.89	165.71	52.92	186.34
n-Pentacosane	519.51	517.29	44.8	526.92
n-Hexacosane	677.44	225.75	111.55	424.36
n-Heptacosane	U	U	26.51	158.31
n-Octacosane	U	U	188.55	1724.06
n-Nonacosane	U	U	U	U
n-Triacontane	U	U	U	U
n-Hentriacontane	U	U	U	U
n-Dotriacontane	U	U	U	U
n-Tritriacontane	U	U	U	U
n-Tetratriacontane	U	U	U	U
n-Pentatriacontane	U	U	U	U
n-Hexatriacontane	U	U	U	U
n-Heptatriacontane	U	U	U	U
n-Octatriacontane	U	U	U	U
n-Nonatriacontane	U	U	U	U
n-Tetracontane	U	U	U	U
TPH(total)	258542.94	306102.65	26740.08	123144.75
Surrogate Recoveries (%)				
O-Terphenyl	107	106	101	111
5a-androstane	96	78	80	91

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07T13	GWP07T14	GWP07S01	GWP07S02
Battelle ID	R5536-P	R5537-P	R5538-P	R5539-P
	SA	SA	SA	
Sample Type				SA
Collection Date	01/11/07	01/11/07	01/12/07	01/12/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/01/07	02/01/07	02/01/07	02/01/07
Analytical Instrument	FID	FID	FID	FID
% Moisture	12.18	7.01	13.85	10.23
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.83	1.95	1.75	1.80
Size Unit-Basis	G_DRY	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit	72.86	2.06	190.48	185.33
Units	MG/KG_DRY	MG/KG_DRY	MG/KG_DRY	MG/KG_DRY
n-Nonane	U	U	U	U
n-Decane	U	U	U	U
n-Undecane	185.62	U	U	U
n-Dodecane	562.64	0.59 J	681.52	345.73
n-Tridecane	2124.98 ME	1.28 JME	806.85 ME	315.85 ME
Isoprenoid RRT 1380	U	U	U	U
n-Tetradecane	154.67	0.55 J	41.31 J	13.81 J
Isoprenoid RRT 1470	174.68	0.37 J	35.44 J	14.1 J
n-Pentadecane	96.34	U	38.51 J	16.02 J
n-Hexadecane	171.28	U	54.63 J	65.43 J
Norpristane (1650)	360.8	U	U	U
n-Heptadecane	66.74 J	0.51 J	14.3 J	U
Pristane	129.3	0.75 J	33.74 J	U
n-Octadecane	2311.44	8.39	3632.75	3464.74
Phytane	90.08	0.48 J	104.55 J	108.04 J
n-Nonadecane	U	U	U	U
n-Eicosane	Ŭ	U	105.71 J	111.97 J
n-Heneicosane	828.22	7.19	3014.6	3804.27
n-Docosane	290.38	7.78	428.84	425.53
n-Tricosane	59.51 J	U	U	U
n-Tetracosane	150.32	7.09	697.9	647.33
n-Pentacosane	67.73 J	4.53	491.26	391.39
n-Hexacosane	240.61	U	263.33	U
n-Heptacosane	U	U	188.48 J	Ŭ
n-Octacosane	U	Ŭ	3959.9	Ŭ
n-Nonacosane	U	U	U	U
n-Triacontane	U	U	U	U
n-Hentriacontane	U	U	Ŭ	U
n-Dotriacontane	U	U	U	U
n-Tritriacontane	U	U	U	U
n-Tetratriacontane	U	U	U	U
n-Pentatriacontane	U	U	U	U
n-Hexatriacontane	U	U	U	U
	U	U	U	U
n-Heptatriacontane				U
n-Octatriacontane	U	U	U	-
n-Nonatriacontane	U	U	U	U
n-Tetracontane	U	U	U	U
TPH(total)	135670.62	2233.63	274222.32	258510.39
Surrogate Recoveries (%)				
O-Terphenyl	98	96	107	110
5a-androstane	84	78	96	108

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07S03	GWP07S04	TDW3-4.5
Battelle ID	R5540-P	R5541-P	R5542-P
Sample Type	SA	SA	SA
Collection Date	01/12/07	01/12/07	09/26/06
Extraction Date	01/24/07	01/24/07	01/24/07
Analysis Date	02/01/07	02/01/07	02/01/07
Analytical Instrument	FID	FID	FID
% Moisture	20.14	12.16	14.86
% Lipid	NA	NA	NA
Matrix	TAR	TAR	SOIL
Sample Size	1.66	1.85	17.17
Size Unit-Basis	G_DRY	G DRY	G DRY
Minimum Reporting Limit	200.8	240.24	1.16
Units	MG/KG_DRY	MG/KG_DRY	MG/KG_DRY
n-Nonane	U	U	UT
n-Decane	U	U	UT
n-Decane n-Undecane	U	U	UT
n-Dodecane	1802	2181.26	0.27 JT
n-Tridecane	9548.63 ME	8148.73 ME	0.27 JT 0.58 JTME
Isoprenoid RRT 1380	9548.63 ME U	0140.73 ME U	UT
n-Tetradecane	1057.6	1071.68	0.18 JT
Isoprenoid RRT 1470	1502.73	1495.76	0.22 JT
n-Pentadecane	418.89	482.53	0.22 JT 0.18 JT
n-Hexadecane	875.13	857.02	0.17 JT
Norpristane (1650)	157.34 J	128.43 J	0.17 JT
n-Heptadecane	674.32	233.72 J	0.31 JT
Pristane	655.85	550.55	0.7 JT
n-Octadecane	3444.29	2423.07	5.9 T
Phytane	118.4 J	187.21 J	0.31 JT
n-Nonadecane	U	U	UT
n-Eicosane	135.2 J	U	UT
n-Heneicosane	1633.15	1599.8	8.69 T
n-Docosane	1117.65	1133.15	4.77 T
n-Tricosane	357.64	81.9 J	0.25 JT
n-Tetracosane	502.47	924.27	9.59 T
n-Pentacosane	371.94	401.16	2.51 T
n-Hexacosane	1360.81	U	7.21 T
n-Heptacosane	U	U	3.72 T
n-Octacosane	U	U	UT
n-Nonacosane	U	U	UT
n-Triacontane	U	U	UT
n-Hentriacontane	U	U	UT
n-Dotriacontane	U	U	UT
n-Tritriacontane	U	U	UT
n-Tetratriacontane	U	U	UT
n-Pentatriacontane	U	U	UT
n-Hexatriacontane	U	U	UT
n-Heptatriacontane	U	U	UT
n-Octatriacontane	U	U	UT
n-Nonatriacontane	U	U	UT
n-Tetracontane	U	U	UT
TPH(total)	421505.26	360563.27	1106.76 T
Surrogate Recoveries (%)			

Surrogate Recoveries (%)

O-Terphenyl	90	115	102
5a-androstane	102	104	80

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	Procedural Blank	
Battelle ID Sample Type Collection Date Extraction Date Analysis Date Analytical Instrument % Moisture % Lipid Matrix Sample Size Size Unit-Basis Minimum Reporting Limit	BJ939PB-P PB 01/24/07 01/24/07 01/31/07 FID 19:52 NA SOIL, TAR 2.44 G_DRY 1.64	
Units	MG/KG_DRY	
n-Nonane n-Decane n-Undecane n-Dodecane n-Tridecane Isoprenoid RRT 1380 n-Tetradecane Isoprenoid RRT 1470 n-Pentadecane n-Hexadecane Norpristane (1650) n-Heptadecane Pristane n-Octadecane Phytane n-Octadecane n-Heneicosane n-Heneicosane n-Heneicosane n-Tricosane n-Tetracosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Heptacosane n-Triacontane n-Triacontane n-Tritriacontane n-Tetratriacontane n-Tetratriacontane n-Tetratriacontane		
n-Hexatriacontane n-Heptatriacontane n-Octatriacontane		U U U
n-Nonatriacontane n-Tetracontane TPH(total)		U U U

Surrogate Recoveries (%)

O-Terphenyl		
5a-androstane		

100 79

The Business of Innovation

Project Client: Exponent, Inc.

Project Name: Exponent - Gas Works Park Project Number: N106746-0001 060208-03: Sand. Client ID White Quartz, -50+70 Battelle ID BJ940LCS-P Sample Type LCS **Collection Date** 01/24/07 Extraction Date 01/24/07 Analysis Date 01/31/07 Analytical Instrument FID % Moisture NA % Lipid NA Matrix SOIL, TAR Sample Size 20.01 Size Unit-Basis G_DRY Minimum Reporting Limit 0.2 MG/KG_DRY Target % Recovery Qualifier Units n-Nonane 0.89 1.25 71 n-Decane 1.25 83 1.04 n-Undecane U n-Dodecane 1.13 1.25 90 n-Tridecane U Isoprenoid RRT 1380 U n-Tetradecane 1.12 1.25 90 Isoprenoid RRT 1470 U n-Pentadecane U n-Hexadecane 1.17 1.25 94 Norpristane (1650) U n-Heptadecane U Pristane 1.25 1.25 100 1.25 n-Octadecane 105 1.31 Phytane 1.15 1.25 92 n-Nonadecane 0.97 1.25 78 n-Eicosane 1.21 1.25 97 U n-Heneicosane n-Docosane 1.24 1.25 99 n-Tricosane U n-Tetracosane 1.21 1.25 97 n-Pentacosane U n-Hexacosane 1.2 1.25 96 n-Heptacosane U n-Octacosane 1.36 1.25 109 n-Nonacosane U n-Triacontane 1.22 1.25 98 n-Hentriacontane U n-Dotriacontane U n-Tritriacontane U n-Tetratriacontane U n-Pentatriacontane U 1.3 n-Hexatriacontane 1.25 104 n-Heptatriacontane U n-Octatriacontane U n-Nonatriacontane U n-Tetracontane U TPH(total) U Surrogate Recoveries (%)

O-Terphenyl 5a-androstane

101

81

The Business of Innovation

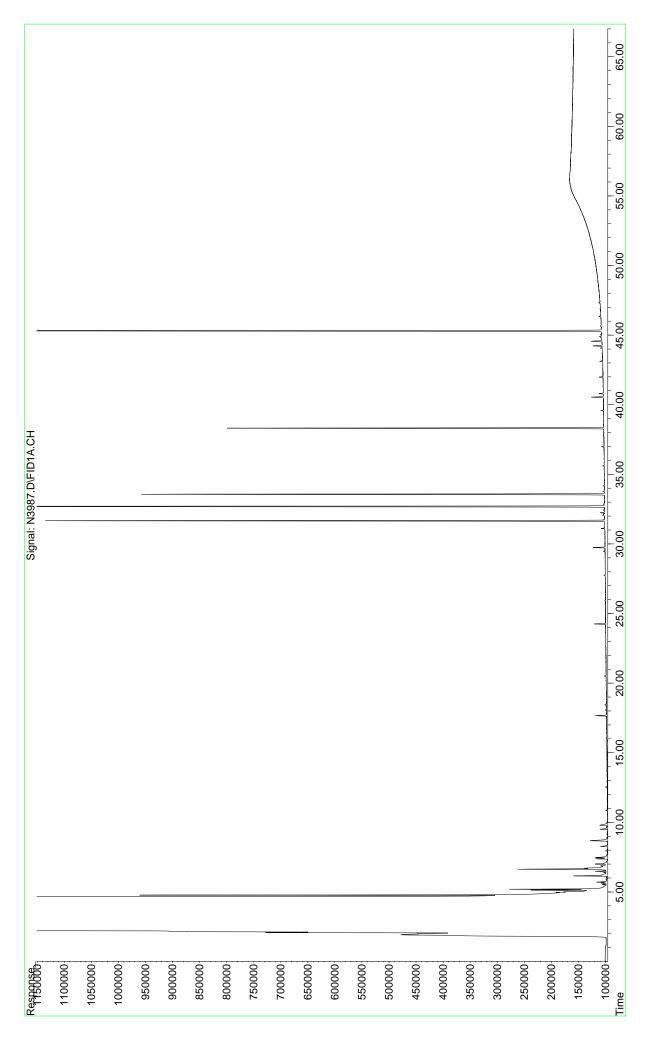
Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GN62: North Slope Crude				
Battelle ID Sample Type	BJ952NSC-P NSC				
Collection Date	01/30/07				
Extraction Date	01/30/07				
Analysis Date	01/31/07				
Analytical Instrument	FID				
% Moisture	NA				
% Lipid	NA				
Matrix	OIL				
Sample Size	5.01				
Size Unit-Basis	MG_OIL				
Minimum Reporting Limit	299.7		Tanatol	D:#	Qualifian
Units	MG/KG_OIL		Target %	Difference	Qualifier
n-Nonane	5020.14		4670.06	7.5	
n-Decane	4863.37		4951.66	1.8	
n-Undecane	4861.8		4506.16	7.9	
n-Dodecane	4899.16		4576.43	7.1	
n-Tridecane	4096.3	ME	4189.33	2.2	
Isoprenoid RRT 1380	965.4		961.81	0.4	
n-Tetradecane	4398.78		3919.50	12.2	
Isoprenoid RRT 1470	1572.82		1532.69	2.6	
n-Pentadecane	4350.22		3990.56	9.0	
n-Hexadecane	3700.26		3640.11	1.7	
Norpristane (1650)	1190.91		1141.72	4.3	
n-Heptadecane	3302.36		3078.38	7.3	
Pristane	2374.46		2280.61	4.1	
n-Octadecane	2877.12		2796.74	2.9	
Phytane	1557.22		1659.88	6.2	
n-Nonadecane	2373.76		2540.37	6.6	
n-Eicosane	2668.31		2502.77	6.6	
n-Heneicosane	2572.46		2419.45	6.3	
n-Docosane	2487.37		2251.79	10.5	
n-Tricosane	2153.59		2050.41	5.0	
n-Tetracosane	2067.1		1948.20	6.1	
n-Pentacosane	1787.61		1795.70	0.5	
n-Hexacosane	1702.6		1639.60	3.8	
n-Heptacosane	1349.47		1230.99	9.6	
n-Octacosane	1105.92		1004.15	10.1	
n-Nonacosane	937.02		872.21	7.4	
n-Triacontane	687.11		669.33	2.7	
n-Hentriacontane	619.57		606.82	2.1 8.2	
n-Dotriacontane	503.97		465.97		
n-Tritriacontane	404.55		399.05 371.75	1.4 1.2	
n-Tetratriacontane n-Pentatriacontane	367.32 383.03		371.75	1.2	
n-Pentatriacontane n-Hexatriacontane	383.03 244.25	J	235.65	1.3 3.6	
n-Heptatriacontane	244.25 214.78		235.65	3.0 2.2	
n-Octatriacontane	214.78 216.26		205.75	2.2 5.1	
n-Nonatriacontane	159.07		153.92	3.3	
n-Tetracontane	172.95	J	161.64	5.5 7.0	
TPH(total)	634437.42	5	578973.63	9.6	
	00		010010.00	5.0	

Surrogate Recoveries (%)

O-Terphenyl	104
5a-androstane	85

File : F:\N\DATA\SN0223\N3987.D Operator : MM Acquired : 1-31-2007 02:46:51 PM using AcqMethod TPH.M Instrument : Inst. N Sample Name: BJ939PB-P-FID(4) Misc Info : Procedural Blank 5-202 07-0010 Vial Number: 6



File : F:\N\DATA\SN0223\N3989.D Operator : MM Acquired : 1-31-2007 04:08:48 PM using AcqMethod TPH.M Instrument : Inst. N Sample Name: BJ940LCS-P-FID(4) Misc Info : Laboratory Control Sample 5-202 07-0010 Vial Number: 7

Response_	-				Signal: N	Signal: N3989.D\FID1A.CH	CH	-				
1050000												
100000												
950000-												
000006												
850000							-					
800000-												
750000												
700000												
650000												
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550000												
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450000												
400000									-			
350000												
30000												
250000												
200000-												
150000												
100000	W.W. W. M. W.							حالالالالك			-	-
Time	5.00 10.00	15.00	20.00	25.00	30.00	35.00	40.00	45.00	50.00	55.00	60.00	65.00

Response	-					Signal: N399	Signal: N3991.D\FID1A.CH	-					
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1800000													
1700000													
1600000													
1500000													
1400000													
1300000			-										
1200000													
1100000													
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-000006					-								
800000													
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600000													
50000													
400000									-				
300000		_								_	-		
200000			ALLONAL AND ALLONAL	ANTALIA, INALIA MUTUTA INA INA	רייזן איזעראווויראין	ANUMUNAD	امتحاط الماريمة المصريا يحدر المحد المصداية الإطارة المالية المالية ومالية المالية المالية المالية المالية الم	u and and and	אההאורייוליולטולטול אוני	hallandur	_ السالسالسا		
100000 ¹ , , , , , , , , , , , , , , , , , , ,	5.00 10.00	0 15.00	0 20.00	-	25.00	30.00	35.00	40.00	45.00	50.00	55.00	60.00	65.00

File :	: F:\N\DATA\SN0223\N3991.D
Operator :	MM
	: 1-31-2007 05:29:00 PM using AcqMethod TPH.M
Ę	Inst. N
Sample Name:	Sample Name: BJ952NSC-P(0)
Misc Info :	: North Slope Crude FID 5-202 07-0010
Vial Number: 8	00

File :		: F:\N\DATA\SN0223\N3993.D
Operator :		MM
	•••	: 1-31-2007 06:49:17 PM using AcqMethod TPH.M
Instrument :		Inst. N
Sample Name:		Sample Name: R5521-P-FID(11)
Misc Info :		Misc Info : GWP07T01 5-202 07-0010
Vial Number: 9		6

Response	_				Signal: N39	Signal: N3993.D/FID1A.CH	Ŧ				
2.8e+07											
2.6e+07											
2.4e+07											
2.2e+07											
2e+07											
1.8e+07-											
1.6e+07-											
1.4e+07-											
1.2e+07-											
1e+07-											
8000000											
600000											
400000					_						
200000							=				
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File :	 : F:\N\DATA\SN0223\N3997.D
Operator :	 MM
Acquired :	 : 1-31-2007 09:29:10 PM using AcqMethod TPH.M
Instrument :	 Inst. N
Sample Name:	 Sample Name: R5525-P-FID(11)
Misc Info :	 Misc Info : GWP07T02 5-202 07-0010
Vial Number: 11	 11

Response_	-				Signal: N35	Signal: N3997.D/FID1A.CH	I					
2.4e+07												
2.2e+07												
2e+07												
1.8e+07												
1.6e+07												
1.4e+07												
1.2e+07												
1e+07												
800000												
600000-												
4000000-												
200000-				-		-						
Time		15 00		**************************************	30.00	35.00	40.00	45 00	50.00	55 00		65.00
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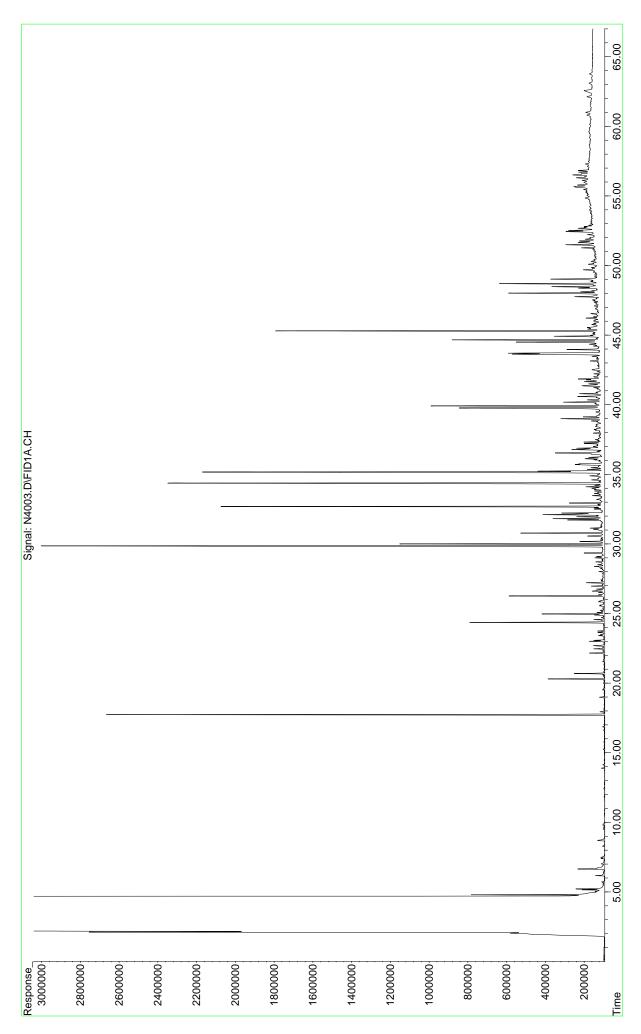
rile	 Бц	: F:\N\DATA\SN0223\N3999.D
Dperator		
Acquired		31 Jan 2007 10:50 pm using AcqMethod TPH.M
Instrument :		Inst. N
Sample Name	 망	Sample Name: R5526-P-FID(11)
disc Info	Ъ.:	1isc Info : GWP07T03 5-202 07-0010
Vial Number: 12		

	- 				

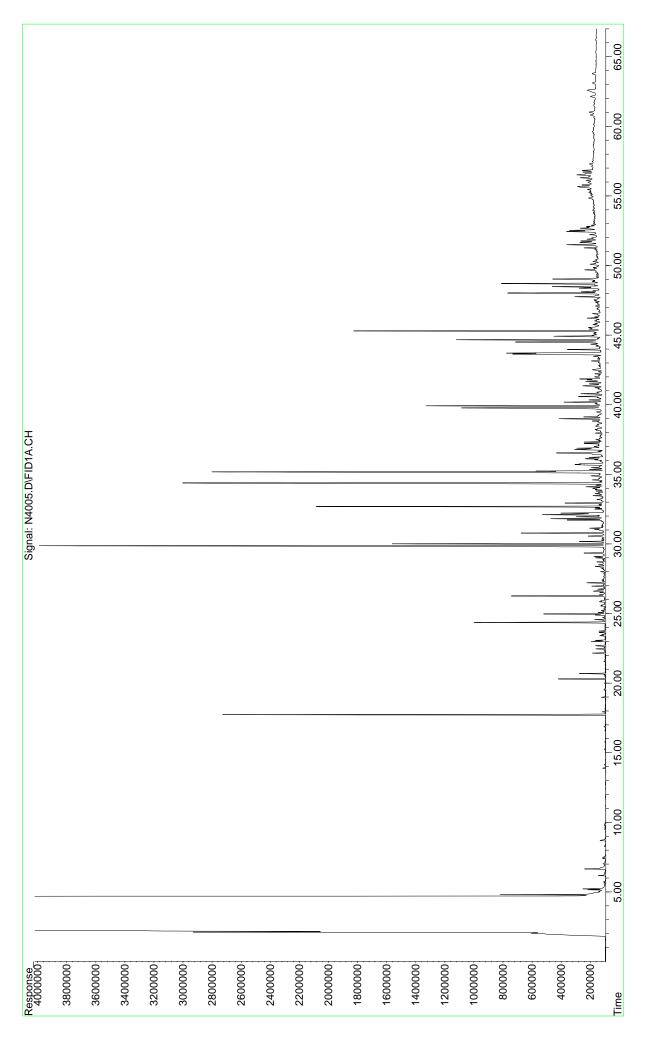
File :	 : F:\N\DATA\SN0223\N4001.D
Operator :	 MM 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1
Acquired :	UL FED ZUU/ 12:11 am USING ACQMETNOG TPH.M
Instrument :	 Inst. N
Sample Name:	 Sample Name: R5527-P-FID(11)
Misc Info :	 Misc Info : GWP07T04 5-202 07-0010
Vial Number: 13	 13

Response2.4e+07				_		Signal: N4	Signal: N4001.D/FID1A.CH	Ŧ				
2.2e+07												
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1.8e+07												
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Time	5.00	10.00	\ .	20.00	<u>₩₩₩₩₩₩₩</u>	30.00	<u> </u>	++	+ \\ \\ 45.00	 55.00	60.00	65.00

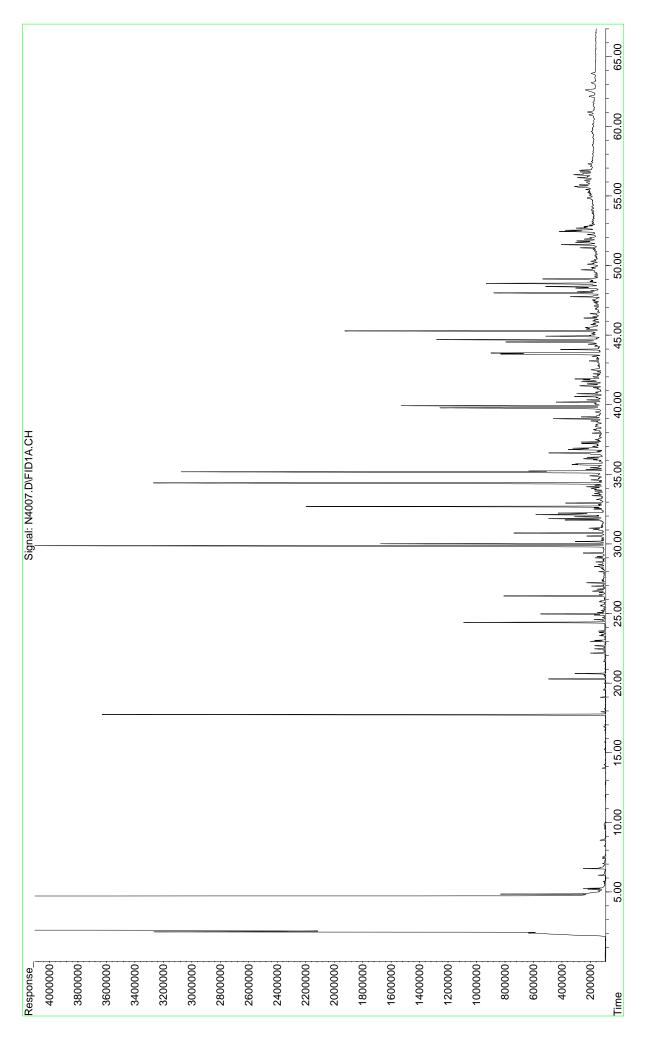
File : F:\N\DATA\SN0223\N4003.D Operator : MM Acquired : 2-1-2007 01:30:53 AM using AcqMethod TPH.M Instrument : Inst. N Sample Name: R5528-P-A-FID(13) Misc Info : GWP07T05 5-202 07-0010 Vial Number: 14



File : F:\N\DATA\SN0223\N4005.D Operator : MM Acquired : 2-1-2007 02:50:25 AM using AcqMethod TPH.M Instrument : Inst. N Sample Name: R5529-P-A-FID(13) Misc Info : GWP07T06 5-202 07-0010 Vial Number: 15



File : F:\N\DATA\SN0223\N4007.D Operator : MM Acquired : 2-1-2007 04:09:54 AM using AcqMethod TPH.M Instrument : Inst. N Sample Name: R5530-P-FID(11) Misc Info : GWP07T07 5-202 07-0010 Vial Number: 16



File :	 : F:\N\DATA\SN0223\N4009.D
Operator :	 MM
Acquired :	 : 2-1-2007 05:29:42 AM using AcqMethod TPH.M
Instrument :	 Inst. N
Sample Name:	 Sample Name: R5531-P-FID(11)
Misc Info :	 Misc Info : GWP07T08 5-202 07-0010
Vial Number: 17	 17

Response			-		Signal: N	Signal: N4009.D/FID1A.CH	H					
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500000-												
4500000												
4000000												
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Time	5.00 10.00	15.00	20.00	1444 Jury Multhum	30.00	1.00 35.00	40.00	45.00	UNULL WWW	55.00	60.00	65.00

File :	F:\N\DATA\SN0223\N4013.D
Operator :	MM
••	2-1-2007 08:09:16 AM using AcqMethod TPH.M
Instrument :	Inst. N
Sample Name:	Sample Name: R5532-P-FID(11)
Misc Info :	GWP07T09 5-202 07-0010
Vial Number: 19	19

Response			_			Signa	al: N4013.	Signal: N4013.D/FID1A.CH	Ŧ						
380000-															
360000															
3400000-															
3200000						-									
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Time	5.00 10.00	0 15.00	-	20.00	25.00	30.00	- 0		40.00	45.00	50.00	55.00	60.00	-	65.00

File :	: F:\N\DATA\SN0223\N4015.D
Operator :	WM
	: 2-1-2007 09:29:43 AM using AcgMethod TPH.M
Ę	Inst. N
Sample Name:	Sample Name: R5533-P-FID(11)
Misc Info :	Misc Info : GWP07T10 5-202 07-0010
Vial Number: 20	20

Response2.1e+07					Signal: N40	Signal: N4015.D/FID1A.CH	Ŧ				
2e+07 1.9e+07											
1.8e+07											
1.7e+07											
1.6e+07											
1.5e+07											
1.4e+07											
1.3e+07											
1.2e+07											
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800000											
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500000		_									
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Lime	10.00	<u>15.00</u>	20.00	25.00	30.00	35.00	40.00	45 00	50.00	55.00	60 00

65.00 60.00 MMM 55.00 50.00 45.00 40.00 Signal: N4017.D\FID1A.CH PW WW WWWW 35.00 UMMMII CANANA MAN 30.00 Acquired:01Feb200710:50 am using AcqMethod TPH.MInstrumentInstrumentInst. NSample Name:R5534-P-FID(11)Misc Info:GWP07T11Vial Number:21 25.00 20.00 15.00 10.00 5.00 2400000 1200000-1000000 800000 600000-400000-200000 1800000-2200000-2000000-1600000 1400000 Response_ Time

: F:\N\DATA\SN0223\N4017.D : MM

Operator Acquired

File

File : F:\N\DATA\SN0223\N4019.D Operator : MM Acquired : 01 Feb 2007 12:10 pm using AcqMethod TPH.M Instrument : Inst. N Sample Name: R5535-P-FID(11) Misc Info : GWP07T12 5-202 07-0010 Vial Number: 22

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Signal: N4019																					+
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Response_	400000	380000	360000	3400000	3200000	300000	280000	260000	2400000	2200000	200000	180000	1600000	1400000	1200000	100000	80000	60000	400000	20000	Time

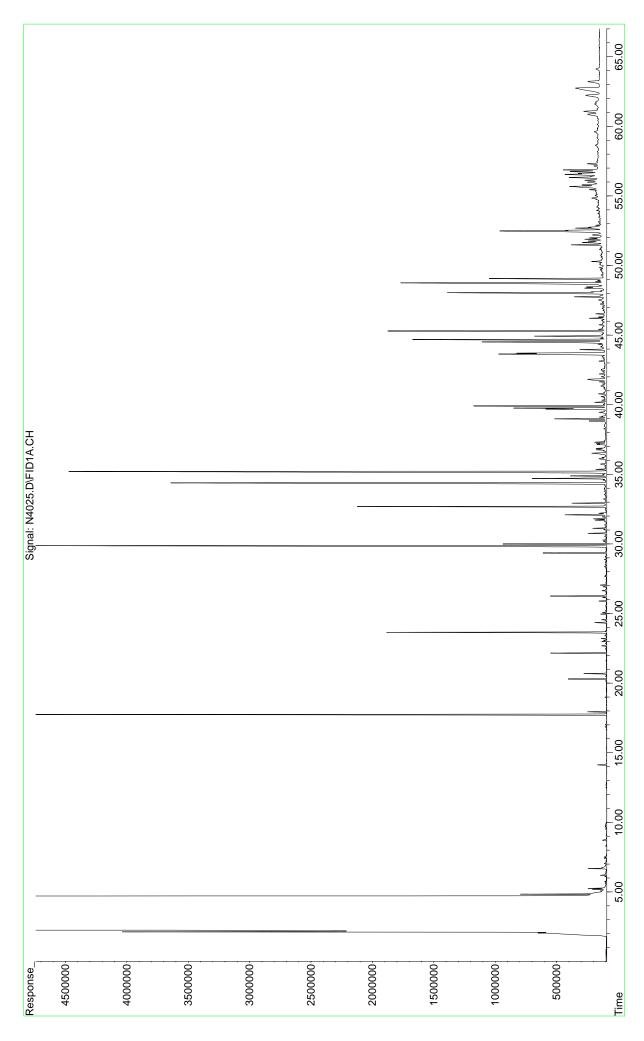
File :	: F:\N\DATA\SN0223\N4021.D
ator	WW
	: 2-1-2007 01:31:24 PM using AcgMethod TPH.M
Ļ	Inst. N
Sample Name:	Sample Name: R5536-P-FID(11)
Misc Info :	Misc Info : GWP07T13 5-202 07-0010
Vial Number: 23	23

Response					U	Signal: N402	Signal: N4021.D\FID1A.CH	Т					
850000													
800000													
750000													
700000													
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500000		-		-	11 11					Munner and Multim	Munum		~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~~
Time	5.00	10.00	15.00	20.00	كالمهممالا	30.00	35.00	40.00	45.00	50.00	55.00	60.00	65.00

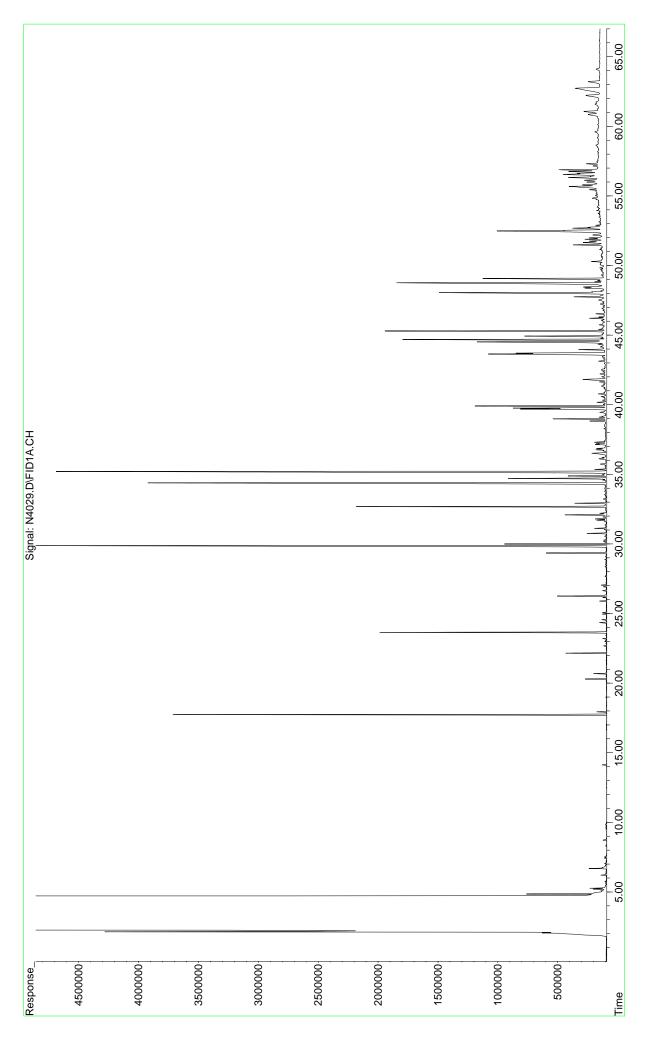
File : F:\N\DATA\SN0223\N4023.D Operator : MM Acquired : 2-1-2007 02:52:05 PM using AcqMethod TPH.M Instrument : Inst. N Sample Name: R5537-P-FID(8) Misc Info : GWP07T14 5-202 07-0010 Vial Number: 24

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File : F:\N\DATA\SN0223\N4025.D Operator : MM Acquired : 2-1-2007 04:12:23 PM using AcqMethod TPH.M Instrument : Inst. N Sample Name: R5538-P-FID(11) Misc Info : GWP07S01 5-202 07-0010 Vial Number: 25



File : F:\N\DATA\SN0223\N4029.D Operator : MM Acquired : 2-1-2007 06:52:57 PM using AcqMethod TPH.M Instrument : Inst. N Sample Name: R5539-P-FID(11) Misc Info : GWP07S02 5-202 07-0010 Vial Number: 27



File		: F:\N\DATA\SN0223\N4031.D
Operator		MM
	•••	: 2-1-2007 08:13:08 PM using AcqMethod TPH.M
Instrument :		Inst. N
Sample Name		Sample Name: R5540-P-FID(11)
Misc Info		Misc Info : GWP07S03 5-202 07-0010
Vial Number: 28	••	28

Response						Signal: N4	Signal: N4031.D/FID1A.CH	A.CH							
6500000															
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	· F · \N \DAIA \SNUZZ3 \N4033.D
Operator :	MM
Acquired :	: 2-1-2007 09:33:07 PM using AcqMethod TPH.M
Instrument :	Inst. N
Sample Name:	Sample Name: R5541-P-FID(11)
Misc Info :	Misc Info : GWP07S04 5-202 07-0010
Vial Number:	29

Response						Signal: N [,]	Signal: N4033.D\FID1A.CH	CH					
5500000													
5000000													
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400000													
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200000						Muldan Lath	Land Marker	Marshall Marshare		Manana ang tang tang tang tang tang tang			
Time	5.00	10.00	15.00	20.00	25.00	30.00	35.00	40.00	45.00	00 30.00 35.00 40.00 45.00 50.00	55.00	60.00	65.00

File : F:\N\DATA\SN0223\N4035.D Operator : MM Acquired : 01 Feb 2007 10:53 pm using AcqMethod TPH.M Instrument : Inst. N Sample Name: R5542-P-A-FID(11) Misc Info : TDW3-4.5 5-202 07-0010 Vial Number: 30

Signal: N4035.D/FID1A.CH									_			0 20.00 25.00 30.00 30.00
											-	4. 1. 10.00 15.00 15.00
		 		 	 	5.00						

SHC and TPH – NAPL QA/QC Summary Batch 07-0011

PROJECT:	Exponent – Gas Works
PARAMETER:	Saturated Hydrocarbons (SHC) and Total Petroleum Hydrocarbons (TPH)
LABORATORY:	Battelle, Duxbury, MA
MATRIX:	Non-aqueous phase liquid (NAPL)
SAMPLE CUSTODY:	Eighteen tar samples, three NAPLs samples, and 1 soil sample were received at the Battelle
	Duxbury Operations (BDO) Laboratory on 1/16/2007. Upon receipt of samples, the
	temperatures of the coolers were taken and the samples were logged into the laboratory and
	given unique IDs. The temperature of the cooler upon receipt was within the acceptable
	range. Samples were either stored in an access-limited walk-in refrigerator at 4°C until
	sample preparation could begin. The NAPL samples were extracted together in one
	analytical batch, batch 07-0011.

QA/QC DATA QUALITY OBJECTIVES:

	Reference Method	Blank	Surrogate Recovery	LCS/MS Recovery	Control Oil % Diff.
SHC and TPH	General NS&T	< 5x MDL	40-120% Recovery	40-120% Recovery MS target spike must be > 5 x background	PD < 30% for 90% of the analytes

METHOD:

NAPL and filter samples were extracted following general NS&T methodologies. Approximately 50 mg of oil was weighed and diluted with 10 mL of hexane, while filter samples were extracted in entirety with 10 mL of Hexane. A portion of the extract was removed and spiked with SIS and IS. One extract was submitted for PAH and petroleum biomarker analysis, and the second extract was submitted for SHC and TPH analysis. NAPL sample data is reported on an oil weight basis, filter sample Bulkhead-02-fp data is reported on a gravimetric weight basis while filter sample Bulkhead Blank data is reported on a ng basis since the sample had no gravimetric weight.

SHC and TPH were measured by gas chromatography with flame ionization detection (GC/FID). An initial calibration consisting of target analytes was completed prior to analysis to demonstrate the linear range of analysis. Calibration verification was performed at the beginning and end of each 12 hour period in which samples were analyzed. Concentrations of SHC and TPH were calculated by the internal standard method. Normal alkanes were quantified using the average RF generated from the initial calibration. TPH concentrations were quantified using the average RF of nC8 through nC40. Isoprenoid hydrocarbon concentrations were quantified using the average RF of the n-alkanes immediately preceding and immediately following each target isoprenoid hydrocarbon.

HOLDING TIMES: Samples were stored cool at approximately 4°C until extraction.

Samples were prepared for analysis in one analytical batch and analyzed within 40 days of extraction.

Batch ID	Extraction Date	Analysis Date(s)
07-0011	1/30/2007	2/5/2007 - 2/6/2007

SHC and TPH – NAPL QA/QC Summary Batch 07-0011

PROCEDURAL BLANK (PB):	A procedural blank (PB) was prepared with each analytical batch. The blank was analyzed to ensure the sample extraction and analysis methods were free of contamination.
	07-0011 – No exceedences noted.
	Comments- None
LABORATORY CONTROL SAMPLE (LCS):	A laboratory control sample (LCS) was prepared each analytical batch. The percent recoveries of target analytes were calculated to measure data quality in terms of accuracy. 07-0011 – No exceedences noted.
	Comments- None
SURROGATE RECOVERY:	Two surrogate compounds were added prior to extraction, including o-terphenyl and 5a- androstane. The recovery of the surrogate compound was calculated to measure data quality in terms of accuracy (extraction efficiency).
	07-0011 – No exceedences noted.
	Comments- None
CONTROL OIL:	A control oil (North Slope Crude) was prepared with the analytical batch. The percent difference (PD) between the measured value and the target value was calculated to measure data quality in terms of accuracy.
	07-0011 – No exceedences noted.
	Comments – None.
CALIBRATIONS:	The GC/FID is calibrated with a minimum 5 level curve for all compounds. The percent relative standard deviation (%RSD) between RF for the individual target analytes must be $\leq 25\%$, and the mean RSD of all target analytes must be $< 20\%$. Each batch of samples analyzed is bracketed by a continuing calibration verification (CCV) sample, run at a frequency of minimally every 12 hours. The PD between the true value and the CCV should be $< 20\%$ for individual analytes. Additionally an initial calibration check (ICC) sample is run immediately after each initial calibration. The PD between the ICC and the initial calibration should be $< 20\%$.
	07-0011 – No exceedences noted.
	Comments – None.

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07DW401	GWP07DW402	GWP07MW9
Battelle ID	R5522-P	R5523-P	R5524-P
Sample Type	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/12/07
Extraction Date	01/30/07	01/30/07	01/30/07
Analysis Date	02/05/07	02/05/07	02/06/07
Analytical Instrument	FID	FID	FID
% Moisture	NA	NA	NA
% Lipid	NA	NA	NA
Matrix	NAPL	NAPL	NAPL
Sample Size	49.50	47.40	45.80
Size Unit-Basis	MG_OIL	MG_OIL	MG_OIL
Reporting Limit	303.03	316.46	327.51
Units	MG/KG_OIL	MG/KG_OIL	MG/KG_OIL
n-Nonane	U	U	U
n-Decane	U	U	U
n-Undecane	985.16	904.4	U
n-Dodecane	2909.61	2982.5	U
n-Tridecane	5232.44 ME	8415.55 ME	4205.06 ME
Isoprenoid RRT 1380	U	U	6331.01
n-Tetradecane	4571.43	4648.67	2968.95
Isoprenoid RRT 1470	3316.27	3374.48	7188.74
n-Pentadecane	1547.03	1667.54	U
n-Hexadecane	1388.3	U	U
Norpristane (1650)	1658.57	363.09	6678.5
n-Heptadecane	946.27	970.61	480.82
Pristane	1377.81	1311.14	10138.83
n-Octadecane	9014.06	9221.92	1509.41
Phytane	317.24	353.6	6173.46
n-Nonadecane	U	U	U
n-Eicosane	468.06	482.61	U
n-Heneicosane	1908.32	1992.63	768.98
n-Docosane	1254.65	1294.69	342.06
n-Tricosane	U	U	U
n-Tetracosane	U	U	U
n-Pentacosane	843.09	868.18	U
n-Hexacosane	1279.06	1262.68	U
n-Heptacosane	205.75 J	224.86 J	U
n-Octacosane	2113.46	2295.97	U
n-Nonacosane	U	U	U
n-Triacontane	U	U	U
n-Hentriacontane	U	U	U
n-Dotriacontane	U	U	U
n-Tritriacontane	U	U	U
n-Tetratriacontane	U	U	U
n-Pentatriacontane	Ū	Ŭ	Ŭ
n-Hexatriacontane	Ŭ	Ŭ	Ŭ
n-Heptatriacontane	U	U	Ŭ
n-Octatriacontane	U	U	U
n-Nonatriacontane	Ŭ	Ŭ	Ŭ
n-Tetracontane	U	U	U
TPH(total)	774165.21	786091.98	869177.39
	777103.21	100091.30	003111.03
Surrogate Recoveries (%)			

O-Terphenyl	107	103	108
5a-androstane	86	87	86

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	Procedural Blank FID	
Battelle ID	BJ963PB-P	
Sample Type	PB	
Collection Date	01/30/07	
Extraction Date	01/30/07	
Analysis Date	02/05/07	
Analytical Instrument	FID	
% Moisture	NA	
% Lipid	NA	
Matrix	OIL	
Sample Size	5.00	
Size Unit-Basis	MG_OIL	
Minimum Reporting Limit	300	
Units	MG/KG_OIL	
Ginto	MG/RG_OL	
n-Nonane		U
n-Decane		U
n-Undecane		U
n-Dodecane		U
n-Tridecane		U
Isoprenoid RRT 1380		U
n-Tetradecane		U
Isoprenoid RRT 1470		U
n-Pentadecane		U
n-Hexadecane		U
Norpristane (1650)		U
n-Heptadecane		U
-		U
Pristane n-Octadecane		U
Phytane		U
n-Nonadecane		U
n-Eicosane		U
		U
n-Heneicosane		U
n-Docosane		U
n-Tricosane		U
n-Tetracosane		
n-Pentacosane		U
n-Hexacosane		U U
n-Heptacosane		U
n-Octacosane		
n-Nonacosane		U U
n-Triacontane		
n-Hentriacontane		U
n-Dotriacontane		U U
n-Tritriacontane		
n-Tetratriacontane		U
n-Pentatriacontane		U
n-Hexatriacontane		U
n-Heptatriacontane		U
n-Octatriacontane		U
n-Nonatriacontane		U
n-Tetracontane		U
TPH(total)		U

Surrogate Recoveries (%)

O-Terphenyl	
5a-androstane	

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	Sample FID				
Battelle ID Sample Type Collection Date Extraction Date	BJ964LCS-P LCS 01/30/07 01/30/07				
Analysis Date	02/05/07				
Analytical Instrument	FID				
% Moisture	NA				
% Lipid Matrix	NA OIL				
Sample Size	NA				
Size Unit-Basis	NA				
Minimum Reporting Limit	1000				
Units	NG		Target	% Recovery	Qualifier
n-Nonane	24417.91		25000.00	98	
n-Decane	25466.5		25000.00	102	
n-Undecane n-Dodecane	25452.29	U	25000.00	102	
n-Tridecane	20402.29	U	23000.00	102	
Isoprenoid RRT 1380		Ŭ			
n-Tetradecane	25215.84		25000.00	101	
Isoprenoid RRT 1470		U			
n-Pentadecane		U			
n-Hexadecane	25548.96		25000.00	102	
Norpristane (1650)		U			
n-Heptadecane Pristane	26973.83	U	25005.00	108	
n-Octadecane	25361.5		25005.00	100	
Phytane	24683.95		25017.50	99	
n-Nonadecane	23835.33		25000.00	95	
n-Eicosane	25536.21		25000.00	102	
n-Heneicosane		U			
n-Docosane	26224.58		25000.00	105	
n-Tricosane	25409.65	U	25000.00	102	
n-Tetracosane n-Pentacosane	25409.65	U	25000.00	102	
n-Hexacosane	25290.62	0	25000.00	101	
n-Heptacosane		U			
n-Octacosane	25240.62		25000.00	101	
n-Nonacosane		U			
n-Triacontane	25174.25		25000.00	101	
n-Hentriacontane		U			
n-Dotriacontane n-Tritriacontane		U U			
n-Tetratriacontane		U			
n-Pentatriacontane		Ū			
n-Hexatriacontane	23884.94		25000.00	96	
n-Heptatriacontane		U			
n-Octatriacontane		U			
n-Nonatriacontane		U			
n-Tetracontane TPH(total)		U U			
i i i i (iotal)		0			

Laboratory Control

Surrogate Recoveries (%)

O-Terphenyl 5a-androstane

99 85

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GN62: North Slope				
Client ID	Crude				
Battelle ID	BJ960NSC-P				
Sample Type	NSC				
Collection Date	01/30/07				
Extraction Date	01/30/07				
Analysis Date	02/05/07				
Analytical Instrument	FID				
% Moisture	NA				
% Lipid	NA				
Matrix	OIL				
Sample Size	5.01				
Size Unit-Basis	MG_OIL				
Minimum Reporting Limit	299.7				
Units	MG/KG_OIL		Target % E	Difference	Qualifier
n-Nonane	5001.37		4670.06	7.1	
n-Decane	4915.31		4951.66	0.7	
n-Undecane	4915.31 4816.79		4506.16	0.7 6.9	
n-Dodecane	4810.79		4576.43	5.5	
n-Tridecane	4726.74	ME	4189.33	12.8	
Isoprenoid RRT 1380	977.58		961.81	1.6	
n-Tetradecane	4166.88		3919.50	6.3	
Isoprenoid RRT 1470	1419.61		1532.69	7.4	
n-Pentadecane	4278.1		3990.56	7.2	
n-Hexadecane	3638.09		3640.11	0.1	
Norpristane (1650)	1145.36		1141.72	0.3	
n-Heptadecane	3308.17		3078.38	7.5	
Pristane	2267.36		2280.61	0.6	
n-Octadecane	2860.32		2796.74	2.3	
Phytane	1535.02		1659.88	7.5	
n-Nonadecane	2453.53		2540.37	3.4	
n-Eicosane	2609.57		2502.77	4.3	
n-Heneicosane	2447.38		2419.45	1.2	
n-Docosane	2466.56		2251.79	9.5	
n-Tricosane	2120.65		2050.41	3.4	
n-Tetracosane	2087.24		1948.20	7.1	
n-Pentacosane	1785.6		1795.70	0.6	
n-Hexacosane	1685.79		1639.60	2.8	
n-Heptacosane	1345.81		1230.99	9.3	
n-Octacosane	1051.9		1004.15	4.8	
n-Nonacosane	868.68		872.21	0.4	
n-Triacontane	668.2		669.33	0.2	
n-Hentriacontane	638.48		606.82	5.2	
n-Dotriacontane	478.41		465.97	2.7	
n-Tritriacontane	389.91		399.05	2.3	
n-Tetratriacontane	355.59		371.75	4.3	
n-Pentatriacontane	360.66		378.11	4.6	
n-Hexatriacontane	234.32	J	235.65	0.6	
n-Heptatriacontane	206.67	J	210.06	1.6	
n-Octatriacontane	218.79	J	205.75	6.3	
n-Nonatriacontane	163.55	J	153.92	6.3	
n-Tetracontane	163.61	J	161.64	1.2	
TPH(total)	612781.39		578973.63	5.8	

Surrogate Recoveries (%)

O-Terphenyl 5a-androstane

98 83

File : Operator : Acquired : Instrument : Sample Name : Misc Info : Vial Number:	: F:\N\DATA\SN0224\N4134.D : MM : 2-5-2007 05:05:46 PM using : Inst. N : BJ963PB-P(0) : Procedural Blank FID 5-202 : 36	\sN0224\ 05:05:46 (0) l Blank	N4134.D PM using FID 5-202	g AcqMethod 2 07-0011	od TPH.M			E					
Kesponse						Signal: N41	Signal: N4134.D/FID1A.CH I	т					
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File : F:\N\DATA\SN0224\N4136.D Operator : MM Acquired : 2-5-2007 06:24:37 PM using AcqMethod TPH.M Instrument : Inst. N Sample Name: BJ964LCS-P(0) Misc Info : Laboratory Control Sample FID 5-202 07-0011 Vial Number: 37

Response 2800000 2600000 2400000 2200000 1800000 1600000 1200000 1000000 800000 600000 400000 800000						Signal: N	Signai: N4136.D/FID1A.CH	D1A.CH							
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Time	5.00	10.00	15.00	20.00	25.00	30.00	35.00		40.00	45.00	50.00	55.00	60.00	65.00	

File	: F:\N\DATA\SN0224\N4138.D
Operator Acquired Instrument : Sample Name: Misc Info : Vial Number:	: MM : 2-5-2007 07:43:23 PM using AcqMethod TPH.M t : Inst. N me: BJ960NSC-P(0) : North Slope Crude FID 5-202 07-0011 er: 38
Response_	Signal: N4138.D/FID1A.CH
1900000	
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File :	: F:\N\DATA\SN0224\N4140.D
<pre>Dperator :</pre>	MM
Acquired :	: 2-5-2007 09:02:02 PM using AcqMethod TPH.M
Instrument :	Inst. N
Sample Name:	Sample Name: R5522-P-FID(5)
Misc Info :	disc Info : GWP07DW401 5-202 07-0011
Vial Number: 39	39

t.2e+07							Signa	II: N4140.I	Signal: N4140.D\FID1A.CH	Ŧ					
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File	: F:\N\DATA\SN0224\N4144.D	
Operator	: MM	
Acquired	: 05 Feb 2007 11:39 pm using AcqMethod TPH.M	
Instrument		
Sample Name	Sample Name: R5523-P-FID(5)	
Misc Info	Misc Info : GWP07DW402 5-202 07-0011	
Vial Number: 41	er: 41	

Response				_			Signi	al: N4144.	Signal: N4144.D\FID1A.CH	ï					
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File : F:\N\DATA\SN0224\N4146.D Operator : MM Acquired : 06 Feb 2007 12:59 am using AcqMethod TPH.M Instrument : Inst. N Sample Name: R5524-P-FID(5) Misc Info : GWP07MW9 5-202 07-0011 Vial Number: 42

Response	_				Signal: N₄	Signal: N4146.D\FID1A.CH	Т					
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Time	5.00 10.00	15.00	20.00	25.00	30.00	35.00	40.00	45.00	50.00	55.00	60.00	65.00

PAH Data and Histograms Biomarker Data and EICPs

PAH and Biomarker – Sediment QA/QC Summary Batches 07-0010

PROJECT:	Exponent – Gas Works Park				
PARAMETER:	Polycyclic Aromatic Hydrocarbons and Biomarkers				
LABORATORY:	Battelle, Duxbury, MA				
MATRIX:	Tar and Soil				
SAMPLE CUSTODY:	Eighteen tar samples, three NAPLs samples, and 1 soil sample were received at the Battelle				
	Duxbury Operations (BDO) Laboratory on 1/16/2007. Upon receipt of samples, the				
	temperatures of the coolers were taken and the samples were logged into the laboratory and				
	given unique IDs. The temperature of the cooler upon receipt was within the acceptable				
	range. Samples were either stored in an access-limited walk-in refrigerator at 4°C until				
	sample preparation could begin. The soil and tar samples were extracted together in one				
	analytical batch, batch 07-0010.				

QA/QC DATA QUALITY OBJECTIVES:

	Reference Method	Blank	Surrogate Recovery	LCS/MS Recovery	MS/MSD Precision	Control Oil % Diff.
PAH and petroleum biomarkers	General NS&T	< 5x MDL	40-120% Recovery	40-120% Recovery MS target spike must be > 5 x background	< 30% RPD	PD < 30% for 90% of the analytes

METHOD:

Soil samples were extracted following general NS&T methodologies. Approximately 15 to 30 grams of soil was spiked with SHC, PAH, and biomarker surrogates and serially extracted three times with dichloromethane using orbital shaker table techniques. The combined extracts were dried over sodium sulfate and concentrated by Kuderna-Danish and nitrogen evaporation techniques. The sample extracts were split in half; one-half of the extract was removed for archiving; the other half was processed through an alumina gravity column to isolate the hydrocarbon fractions of interest. The weight of the resulting extract was determined gravimetrically. The extracts were concentrated to 1 ml, split, and spiked with IS. The pre-injection volume and/or extract split were adjusted to achieve 3 mg/mL. One extract was submitted for PAH and petroleum biomarker analysis, and the second extract was submitted for SHC and TPH analysis.

PAH and petroleum biomarkers were measured by gas chromatography-mass spectrometry (GC/MS) in the selected ion mode (SIM). An initial calibration consisting of target analytes was analyzed prior to analysis to demonstrate the linear range of analysis. Calibration verification was performed at the beginning and end of each 12 hour period in which samples were analyzed. Concentrations of PAH and petroleum biomarkers were calculated versus internal standards. Target PAH were quantified using the average response factors (RF) generated from the initial calibration. The alkyl homolgue PAH series were assigned the RF of the parent PAH, steranes were assigned the RF of cholestane, and triterpanes were assigned the RF of moretane.

Note: the reporting limit for alkylbenzene compounds is orders of magnitude higher than the reporting limit for the rest of the PAH compounds.

PAH and Biomarker – Sediment QA/QC Summary Batches 07-0010

HOLDING TIMES:	14 days of sample receipt collection to extraction fo the storage conditions for	and analyzed within 40 day r soil samples are 14-days i	batch. Samples were extracted within ys of extraction. Holding times from f refrigerated, 365-days if frozen. Since ed 9/26/2006, prior to its arrival at BDO fied with a "T".			
	Batch ID 07-0010	Extraction Date 1/24/2007	Analysis Date(s) 2/7/2006 – 2/14/2007			
PROCEDURAL BLANK (PB):		was prepared with each ana on and analysis methods w	lytical batch. The blank was analyzed to ere free of contamination.			
	07-0010 – 1 exceedence m	oted.				
	the MDL (8.42 ng/g). Fie		at a concentration greater than five times or naphthalene were significantly higher rective action required.			
LABORATORY CONTROL SAMPLE (LCS):			h analytical batch. The percent ure data quality in terms of accuracy.			
	07-0010 - No exceedence	es noted.				
	Comments – None.					
SURROGATE RECOVERY:	Five surrogate compounds were added prior to extraction, including d8-naphthalene, d10- acenaphthene, d10-phenanthrene, and d12-benzo(a)pyrene, and 5(b)H-cholane. The recovery of the surrogate compound was calculated to measure data quality in terms of accuracy (extraction efficiency).					
	07-0010 – 31 exceedences noted.					
	Comments – (b)H-cholane was over-recovered in all field samples. d8-Naphthalene was under-recovered in samples GWP07T01and GWP07T03. d-12 Benzo(a)pyrene was over-recovered in samples GWP07T05, GWP07T07, GWP07T08, GWP07T09, GWP07T12, GWP07T13, GW07S01, GWP07S02, GWP07S03, and GWP07S04. All surrogate exceedences are due to the high contamination level in the samples, and have been qualified with an "NME" to indicate the exceedences are due to matrix effect.					
CONTROL OIL:		he measured value and the	the analytical batch. The percent target value was calculated to measure			
	07-0010 – No exceedence	es noted.				
	Comments – None.					

PAH and Biomarker – Sediment QA/QC Summary Batches 07-0010

CALIBRATIONS:The GC/MS is calibrated with a minimum 5 level curve for all compounds. The percent
relative standard deviation (%RSD) between RF for the individual target analytes must be
 \leq 30%, and the mean RSD of all target analytes must be <15%. Each batch of samples
analyzed is bracketed by a continuing calibration verification (CCV) sample, run at a
frequency of minimally every 12 hours. The PD between the true value and the CCV should
be <25% for individual analytes. Additionally an initial calibration check (ICC) sample is run
immediately after each initial calibration. The PD between the ICC and the initial calibration
should be <25%.</td>

07-0010 - No exceedences noted.

Comments – None.

Battelle

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07T01	GWP07T02	GWP07T03	GWP07T04
Battelle ID	R5521-P	R5525-P	R5526-P	R5527-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/11/07	01/11/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/12/07	02/10/07	02/09/07	02/09/07
Analytical Instrument	MS	MS	MS	MS
% Moisture	28.88	50.1	50.3	34.93
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.44	1.02	1.01	1.44
Size Unit-Basis	G_DRY	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit Units	281.14 NG/G_DRY	1240.34 NG/G_DRY	1044.29 NG/G_DRY	488.3 NG/G_DRY
Onits	NG/G_DK1	NG/G_DR1	NG/G_DK1	NO/O_DKT
C3-Alkylbenzenes	200241.89	740500.51	544602.3	320855.54
C4-Alkylbenzenes	127384.2	392300.28	291736.71	199914.12
C5-Alkylbenzenes	23379.2 J	69444.5 J	51860.33 J	37371.72 J
C6-Alkylbenzenes	9519.52 J	26200.83 J	19546.42 J	14739.7 J
Benzo(b)thiophene	808170.83	2934365.74	2091464.31	1072137.6
C1-benzo(b)thiophenes	82288.56	219469.33	161209.05	118049
C2-benzo(b)thiophenes	25587.92	63190.68	45916.09	34531.23
C3-benzo(b)thiophenes	6724.18	17958.79	13558.37	8978.88
C4-benzo(b)thiophenes	1516.3	4177.36	2615.75	1908.26
Naphthalene	87981028.34 D 2197272.8	378914338.6 D	248684663.2 D 4544807	113638067.2 D 3162312.68
C1-Naphthalenes C2-Naphthalenes	599102.14	6076038.76 1518395.61	1092221.59	817449.67
C3-Naphthalenes	197880.46	477415.52	336362.53	256499.08
C4-Naphthalenes	52064.86	127870.43	85971.42	65429.99
Biphenyl	454683.59	1184039.28	868480.2	639554.02
C1-Biphenyls + Dibenzofuran	966212.61	2460541.43	1791369.85	1328068.78
C2-Biphenyls + C1-Dibenzofurans	262579.31	637542.98	455780.84	349172.44
Acenaphthylene	1973285.53 D	5537642.35 D	4060515.93 D	2692348.15 D
Acenaphthene	285034.92	709485.38	510151.04	375293.53
Dibenzofuran	837673.49	2145426.95	1564653.17	1156883.78
Fluorene	904621.84	2349835.85	1683830.72	1205960.13
C1-Fluorenes	112323.51	292247.97	179830.74	129613.09
C2-Fluorenes	41956.52	90599.94	63677.05	45832.25
C3-Fluorenes	28090.04	55225.82	43056.81	41673.07
C4-Fluorenes Anthracene	23282.04 748435.91 D	37973.81 2660367.97	38625 1900732.77	29355.34 1340503.68
Phenanthrene	3719663.76 D	8334306.16 D	6784198.36 D	5438479.44 D
C1-Phenanthrenes/Anthracenes	545149.65	1316736.52	922450.96	709667.29
C2-Phenanthrenes/Anthracenes	175754.19	384891.14	280592.34	209185.93
C3-Phenanthrenes/Anthracenes	60038.95	137878.3	96256.35	71843.81
C4-Phenanthrenes/Anthracenes	16755.98	37901.3	30011.68	20688.25
Retene	16608.63	37693.1	25082.64	18792.64
Dibenzothiophene	156443.91	386954.46	273906.09	208212.9
C1-Dibenzothiophenes	31741.48	77811.54	53431.96	40801.72
C2-Dibenzothiophenes	11763.09	28766.57	19936.93	15567.12
C3-Dibenzothiophenes	5766.46	13019.78	9164.7	6731.11
C4-Dibenzothiophenes	1556.21	5215.75	2668.13	1791.68
Fluoranthene	2667711.89 D	6197361.74 D	4646671.1 D	3690165.6 D
Pyrene	2277073.94 D	4820682.77 D	3900598.32 D	3063545.13 D
C1-Fluoranthenes/Pyrenes	604569.61	1510567.23	1044364.86	740920.57
C2-Fluoranthenes/Pyrenes	157177.94	324998.68	233225.34	193823.39
C3-Fluoranthenes/Pyrenes C4-Fluoranthenes/Pyrenes	49635.71 27127.96	124029.31 38210.82	83739.63 48389.03	66622.04 35783.93
C0-Benzo(b)naphthothiophenes	39868.25	93925.17	66452.3	52190.34
C1-Benzo(b)naphthothiophenes	18780.12	42078.27	27938.49	23608.21
C2-Benzo(b)naphthothiophenes	5816.1	13668.57	10299.92	6895.74
C3-Benzo(b)naphthothiophenes	7116.14	16044.83	13620.22	8640.99
C4-Benzo(b)naphthothiophenes	1267.71	3337.31	1625.15	1836.15
Benzo(a)anthracene	779123.01	1881479.4	1329724.3	991494.48
Chrysene	735216.55	1705008.49	1235623.91	919283.01

Battelle

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07T01	GWP07T02	GWP07T03	GWP07T04
Battelle ID	R5521-P	R5525-P	R5526-P	R5527-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/11/07	01/11/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/12/07	02/10/07	02/09/07	02/09/07
Analytical Instrument	MS	MS	MS	MS
% Moisture	28.88	50.1	50.3	34.93
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.44	1.02	1.01	1.44
Size Unit-Basis	G_DRY	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit	281.14	1240.34	1044.29	488.3
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
C1-Chrysenes	186899	427281.94	293837.11	223996.13
C2-Chrysenes	63142.39	146843.07	99966.78	72766.81
C3-Chrysenes	32332.16	106991.93	80682.88	49488.17
C4-Chrysenes	8735.48	14005.92	12381.17	8380.26
Benzo(b)fluoranthene	613443.59	1298687.15	949836.19	752475.2
Benzo(k)fluoranthene	695610.31	1645954.33	1155474.88	866874.81
Benzo(e)pyrene	554569.65	1255137.45	911100.65	689339.59
Benzo(a)pyrene	1065303.82	2408803.35	1719568.32	1289502.51
Pervlene	345127.64	740509.08	528443.28	410475.45
Indeno(1,2,3-cd)pyrene	761702.38	1731032.79	1240056.85	961042.01
Dibenz(a,h)anthracene	119887.72	240723.05	168280.12	131715.72
Benzo(g,h,i)perylene	764495.1	1720045.71	1254098.16	965716.53
Total PAH	113619845.5	441162555.3	294979245.5	148378016.9
Surrogate Recoveries (%)				
Naphthalene-d8	24 NME	70	38 NME	67
Acenaphthene-d10	73	78	66	70
Phenanthrene-d10	80	90	80	87
Benzo(a)pyrene-d12	110	96	94	94

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Client ID	GWP07T05	GWP07T06	GWP07T07	GWP07T08
Battelle ID	R5528-P	R5529-P	R5530-P	R5531-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/11/07	01/11/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/08/07	02/08/07	02/08/07	02/11/07
Analytical Instrument	MS	MS	MS	MS
% Moisture	4.23	6.94	6.47	30.28
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.92	1.93	1.87	1.49
Size Unit-Basis	G_DRY	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit	1449.65	969.51	992.27	934.01
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
C3-Alkylbenzenes	16643.82 J	11632.94 J	11720.98 J	35938.6 J
C4-Alkylbenzenes	18269.2 J	13375.85 J	13845.59 J	65757.51 J
C5-Alkylbenzenes	4900.61 J	3514.58 J	3497.96 J	12543.17 J
C6-Alkylbenzenes	1894.64 J	U	982.99 J	7874.5 J
Benzo(b)thiophene	170508.78	108387.93	141819.22	782246.38
C1-benzo(b)thiophenes	43922.65	31332.87	35349.83	605540.6
C2-benzo(b)thiophenes	30116.14	24141.77	25158.21	549105.87
C3-benzo(b)thiophenes	13843.29	12339.22	11883.37	177168.13
C4-benzo(b)thiophenes	4725.35	4270.95	3571.41	29031.26
Naphthalene	12799972.26 D	8370861.88 D	11298733.16 D	22267654.71 D
C1-Naphthalenes	1486915.45	1078913.71	1185207.88	6376462.97
C2-Naphthalenes	884717.78	731690.04	732986.71	3933479.51
C3-Naphthalenes	465601.45	427265.02	413581.2	1319799.8
C4-Naphthalenes	186182.16	182598.98	173957.22	234321.2
Biphenyl	397569.91	304501.81	321153.01	1370918.67
C1-Biphenyls + Dibenzofuran	2422635.2	2080320.47	1967361.36	757779.53
C2-Biphenyls + C1-Dibenzofurans	959968.41	896783.56	844367.87	509842.78
Acenaphthylene	44864.09	31802.1	28696.3	2408513.29
Acenaphthene	3770326.74	3018808.59 D	3283019.88 D	316370.51
Dibenzofuran	2066229.18	1775620.25	1782695.2 D	371781.72
Fluorene	2673369.51	2279642.43	2343495.76 D	2210941.52
C1-Fluorenes	432142.76	403061	411620.05	747906.54
C2-Fluorenes	200826.92	190577.73	222271.76	255951.1
C3-Fluorenes	161247.38	200254.06	171384.74	157418.35
C4-Fluorenes	146309.67	145878.09	152394.9	61709.87
Anthracene	5353451.21 D	5197306.26 D	5324002.63 D	2576612.15
Phenanthrene C1-Phenanthrenes/Anthracenes	16503180.16 D	15593447.41 D	16283960.62 D	13778166.28 D
C1-Phenanthrenes/Anthracenes	3738000.72	3513357.24 1415337.3	3369621.5 1439871.86	3777576.77 1177836.63
C3-Phenanthrenes/Anthracenes	1460972.93 604353.09	576273.77	603675.97	254939.68
C4-Phenanthrenes/Anthracenes	169873.5	170503.18	201299.14	40945.67
Retene	351280.25	345423.48	359624.96	40945.67 U
Dibenzothiophene	562702.85	526970.55	498078.59	1306618.92
C1-Dibenzothiophenes	150631.44	145321.6	142086.32	611469.7
C2-Dibenzothiophenes	64336.15	63060.51	64238.87	292654.59
C3-Dibenzothiophenes	32726.6	30226.36	32725.71	87040.51
C4-Dibenzothiophenes	9836.09	7614.17	11307.3	14206.06
Fluoranthene	12427734.3 D	11851310.8 D	12984354.54 D	6410742.12 D
Pyrene	12257379.44 D	11684717.46 D	12912726.03 D	8576504.35 D
C1-Fluoranthenes/Pyrenes	3666549.68	3248534.17	3512515.4	2565017.43
C2-Fluoranthenes/Pyrenes	958652.35	965239.01	1056810.25	736260.09
C3-Fluoranthenes/Pyrenes	396583.89	438139.89	436357.63	194956.72
C4-Fluoranthenes/Pyrenes	183519.7	193563.04	213915.67	46774.56
C0-Benzo(b)naphthothiophenes	202024.03	188707.18	200086.12	351643.5
C1-Benzo(b)naphthothiophenes	99858.18	100235.56	103812.64	250739.06
C2-Benzo(b)naphthothiophenes	36081.65	32482.84	39025.68	105906.27
C3-Benzo(b)naphthothiophenes	39767.85	40157.99	47446.73	59034.8
C4-Benzo(b)naphthothiophenes	10500.81	11174.56	14645.47	14094.95
Benzo(a)anthracene	3848884.26	3691087.79 D	4139655.69 D	2121727.42
Chrysene	3664745.85	3625200.38 D	4057987.38 D	2437418.12

The Business of Innovation

Client ID	GWP07T05	GWP07T06	GWP07T07	GWP07T08
Battelle ID	R5528-P	R5529-P	R5530-P	R5531-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/11/07	01/11/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/08/07	02/08/07	02/08/07	02/11/07
Analytical Instrument	MS	MS	MS	MS
% Moisture	4.23	6.94	6.47	30.28
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.92	1.93	1.87	1.49
Size Unit-Basis	G_DRY	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit	1449.65	969.51	992.27	934.01
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
C1-Chrysenes	1148640.89	1073725.91	1212316.18	1003941.42
C2-Chrysenes	428957.15	436161.25	535767.32	361014.6
C2-Chrysenes	303677.61	281809.93	338022.76	150152.43
C4-Chrysenes	45363.1	56221.48	84118.52	34853.32
Benzo(b)fluoranthene	2651645.19	2613760.42	2722647.73 D	1530971.59
Benzo(k)fluoranthene	3316052.5	3043902.3	3009453.47	1914736.39
Benzo(e)pyrene	2633480.75	2503298.29	2697597.76 D	1774418.3
Benzo(a)pyrene	4643724.57 D	4252114.79 D	4862823.78 D	3113342.2
Perylene	1417611.53	1367073.07	1580992.44	812754.07
Indeno(1,2,3-cd)pyrene	3311073.71	2792696.64 D	3189174.8 D	2288335.56
Dibenz(a,h)anthracene	503905.76	534705.23	605991.98	386713.77
Benzo(g,h,i)perylene	3245118.71	3000716.48 D	3400513.54 D	2220649.95
Total PAH	115089811.6	103695431.2	113679498.6	104524096.7
Surrogate Recoveries (%)				
Naphthalene-d8	88	104	101	81
Acenaphthene-d10	88	79	81	68
Phenanthrene-d10	88	94	87	77
Benzo(a)pyrene-d12	399 NME	89	411 NME	157 NME

The Business of Innovation

Client ID	GWP07T09	GWP07T10	GWP07T11	GWP07T12
Battelle ID	R5532-P	R5533-P	R5534-P	R5535-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/11/07	01/11/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/08/07	02/09/07	02/11/07	02/08/07
Analytical Instrument	MS	MS	MS	MS
% Moisture	19.46	32.51	5.51	10.85
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.72	1.45	2.05	1.79
Size Unit-Basis	G DRY	G_DRY	G DRY	G_DRY
Minimum Reporting Limit	1078.81	698.01	977.56	414.65
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
		_		_
C3-Alkylbenzenes	36580.22 J	525029.89	2191.57 DJ	39680.66 J
C4-Alkylbenzenes	71566.13 J	267770.56	5055.11 DJ	44626.03 J
C5-Alkylbenzenes	13353.4 J	50020.43 J	1590.34 DJ	7993.18 J
C6-Alkylbenzenes	9286.1 J	18809.7 J	1132.86 DJ	4237.2 J
Benzo(b)thiophene	600997.35	1409434.13	6602.21 D	202487.18
C1-benzo(b)thiophenes	559175.82	138298.11	18195.82 D	82126.5
C2-benzo(b)thiophenes	580029.73	40696.8	32674.4 D	36340.56
C3-benzo(b)thiophenes	207713.37	11562.9	20464.75 D	9868.5
C4-benzo(b)thiophenes	36712.3	2649.01	6076.18 D	1944.66
Naphthalene	15284889.93 D	145298242.7 D	223304.71 D	4999828.2 D
C1-Naphthalenes	6211078.04	3899752.92	350693.91 D	2205708.83
C2-Naphthalenes	4284353.22	970727.61	352583.57 D	809484.01
C3-Naphthalenes	1569159.41	306810.83	137220.32 D	248499.46
C4-Naphthalenes	293313.36	82897.14	36593.43 D	64641.58
Biphenyl	1353389.83	753471.08	50124.82 D	317396
C1-Biphenyls + Dibenzofuran	814568.89	1566453.23	103159.32 D	1263797.88
C2-Biphenyls + C1-Dibenzofurans	617126.67	410565.66	84113.06 D	478999.88
Acenaphthylene	2723434.28	3466032.62 D	70113.9 D	1849540.36 D
Acenaphthene	322276.01	444398.4	238718.81 D	266858.11
Dibenzofuran	381628.07	1361607.06	61672.29 D	1201966.66 D
Fluorene	2430112.88	1491341.86	245201.44 D	1473457.49 D
C1-Fluorenes	926137.56	178262.98	113263.88 D	194027.96
C2-Fluorenes	355186.83	66723.22	51536.29 D	73084.62
C3-Fluorenes	179920.17	37615.8	18702.8 D	85391.42
C4-Fluorenes	76583.75	40720.93	12491.92 D	51229.37
Anthracene	2965638.91	1750201.39	277290.96 D	1620907.51 D
Phenanthrene	14428616.22 D	7263752.38 D	1060654.5 D	6996121.82 D
C1-Phenanthrenes/Anthracenes	4511051.53	862740.73	500955.37 D	1230217.63
C2-Phenanthrenes/Anthracenes	1474202.26	278082.25	179938.85 D	433471.24
C3-Phenanthrenes/Anthracenes	330344.99	97771.75	48951.4 D	136666.94
C4-Phenanthrenes/Anthracenes	57750.41	28485.65	11119.04 D	38936.33
Retene	U	36778.51	4865.52 D	52945.19
Dibenzothiophene	1446533.18	247820.26	79816.35 D	257089.58
C1-Dibenzothiophenes	732148.82	49989.6	74607.4 D	65380.27
C2-Dibenzothiophenes	365443.44	19653.69	45202.6 D	25930.43
C3-Dibenzothiophenes	116396.7	8170.17	17548.54 D	10655.43
C4-Dibenzothiophenes	21781.27	2639.63	4933.51 D	2258.07
Fluoranthene	7291103.73 D	5056902.67 D	519719.5 D	5371309.03 D
Pyrene	10065014.19 D	4262558.61 D	632498.36 D	5122420.76 D
C1-Fluoranthenes/Pyrenes	3272368.71	1009297.21	374611.91 D	1270064.14
C2-Fluoranthenes/Pyrenes	937168.77	229850.65	110112.61 D	427613.13
C3-Fluoranthenes/Pyrenes	246295.46	80465.28	31944.82 D	125298.55
C4-Fluoranthenes/Pyrenes	62759.66	49537.86	8622.3 D	53036.15
C0-Benzo(b)naphthothiophenes	423790.53	65863.38	27427.54 D	115923.36
C1-Benzo(b)naphthothiophenes	302029.96	30065.31	27932.63 D	52530.21
C2-Benzo(b)naphthothiophenes	131066.21	10891.34	12159.6 D	16868.85
C3-Benzo(b)naphthothiophenes	77214.29	11772.33	5496.42 D	16075.47
C4-Benzo(b)naphthothiophenes	16587.38	2259.44	2178.97 D	2823.57
Benzo(a)anthracene	2488334.87	1273089.82	228885.55 D	1642711.55 D
Chrysene	2867991.71	1190069.72	236434.16 D	1614818.64 D

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Client ID	GWP07T09	GWP07T10	GWP07T11	GWP07T12
Battelle ID	R5532-P	R5533-P	R5534-P	R5535-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/11/07	01/11/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/08/07	02/09/07	02/11/07	02/08/07
Analytical Instrument	MS	MS	MS	MS
% Moisture	19.46	32.51	5.51	10.85
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.72	1.45	2.05	1.79
Size Unit-Basis	G_DRY	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit	1078.81	698.01	977.56	414.65
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
C1-Chrysenes	1215740.74	298416.09	116117.61 D	430987.26
C2-Chrysenes	425207.68	97163.86	33587.03 D	145186.25
C3-Chrysenes	158003.22	72744.66	11370.82 D	70869.11
C3-Chrysenes C4-Chrysenes	41669.54	10381.95	2119.89 D	18003.21
Benzo(b)fluoranthene	1807708.96	965211.2	102703.46 D	1193235.69 D
Benzo(k)fluoranthene	2216835.94	1146342.11	145824.59 D	1441182.67
Benzo(e)pyrene	2048385.44	913574.37	143824.39 D 113381.94 D	1135788.54 D
Benzo(a)pyrene	3686081.03	1736642.79	216113.91 D	1969951.25 D
Perylene	941296.51	547303.45	42255.27 D	698839.98
Indeno(1,2,3-cd)pyrene	2601656.25	1220562.11	42255.27 D 108369.76 D	1488582.15 D
Dibenz(a,h)anthracene	432410.31	170838	24908.62 D	301431.86
Benzo(g,h,i)perylene	2477511.48	1266127.39	104107.18 D	1620078.65 D
Total PAH	107985571.9	190514733.7	7405815.68	50695892.37
Idaran	107903371.9	190314733.7	7403013.00	30093092.37
Surrogate Recoveries (%)				
Naphthalene-d8	99	41	78 D	94
Acenaphthene-d10	88	56	54 D	61
Phenanthrene-d10	84	83	80 D	89
Benzo(a)pyrene-d12	194 NME	111	89 D	138 NME

The Business of Innovation

Client ID	GWP07T13	GWP07T14	GWP07S01	GWP07S02
Battelle ID	R5536-P	R5537-P	R5538-P	R5539-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/12/07	01/12/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/09/07	02/09/07	02/09/07	02/08/07
Analytical Instrument	MS	MS	MS	MS
% Moisture	12.18	7.01	13.85	10.23
% Lipid	NĂ	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.83	1.95	1.75	1.80
Size Unit-Basis	G DRY	G DRY	G DRY	G_DRY
Minimum Reporting Limit	384.24	12.91	1004.51	1030.86
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
C3-Alkylbenzenes	190546.76	42.74 J	9764.91 J	2246.68 J
C4-Alkylbenzenes	125716.17	29.24 J	8066.24 J	1603.81 J
C5-Alkylbenzenes	20333.3 J	19.14 J	2046.11 J	U
C6-Alkylbenzenes	8708.41 J	16.89 J	1470.33 J	U
Benzo(b)thiophene	563452.05	292.66	758954.67	395927.35
C1-benzo(b)thiophenes	131790.2	150.38	86699.74	36079.64
C2-benzo(b)thiophenes	41945.85	221.93	42898	13843.85
C3-benzo(b)thiophenes	9133.19	170.14	14447.97	3948.09
C4-benzo(b)thiophenes	1706.06	115.18	3549.83	897.61 J
Naphthalene	16263103.13 D	5059.57	21721142 D	12234462.62 D
C1-Naphthalenes	3314100.42	1481.53	1127445.06	623848.67
C2-Naphthalenes	905688.15	1564.22	246488.39	84945.32
C3-Naphthalenes	268147.15	1130.83	64876.2	18541.83
C4-Naphthalenes	87700.42	778.54	15114.43	4413.73
Biphenyl	632479.48	995.72	1562671.63	1113379.92
C1-Biphenyls + Dibenzofuran	1524802.87	590.82	291733.82	192543.94
C2-Biphenyls + C1-Dibenzofurans	482761.56	646.64	57689.31	34707.76
Acenaphthylene	2847800.61 D	3922.19	7231603.33 D	6490099.33 D
Acenaphthene	271355.89	187.75	232629.82	112631.05
Dibenzofuran	1652087.79 D	322.84	206558.66	144056.33
Fluorene	1725178.93 D	777.53	1676012.74	1438195.8
C1-Fluorenes	181489.89	568.29	102492.82	57887.68
C2-Fluorenes	64836.33	1136.75	29679.66	14011.39
C3-Fluorenes	66157.24	2655.14	23834.33	10175.46
C4-Fluorenes	44910.49	3959.41	7772.65	3654.53
Anthracene	1607520.18 D	4195.11	2774651.05	2925100.01
Phenanthrene	8192705.83 D	21141.12	28059893.4 D	21867409.72 D
C1-Phenanthrenes/Anthracenes	1083410.58	8694.55	939016.67	770315.86
C2-Phenanthrenes/Anthracenes	367874.88	11122.07	151458.42	88878.02
C3-Phenanthrenes/Anthracenes	117746.98	7938.89	42447.12	22664.61
C4-Phenanthrenes/Anthracenes	35588.9	2817.46	7548.84	3506.41
Retene	55643.33	726.77	2239.34	845.44 J
Dibenzothiophene	253540.14	2104.04	2296634.96	2175735.95
C1-Dibenzothiophenes	57973.86	1438.73 2547.28	141231.41	101150.11 23842.63
C2-Dibenzothiophenes	19815.42		42580.56	
C3-Dibenzothiophenes	8012.08	2413.97	18234.26	9336.34 2196.64
C4-Dibenzothiophenes Fluoranthene	1692.49 5676062.55 D	1000.78 39773.81	4058.68 21361152.11 D	2196.64 17772525.52 D
Pyrene	5293384.47 D	65047.13 D	29709883.05 D	23700020.48 D
C1-Fluoranthenes/Pyrenes	1087088.92	22213.95	1927284.7	1862567.79
C2-Fluoranthenes/Pyrenes	317097.28	15552.37	195657.49	148188.86
C3-Fluoranthenes/Pyrenes	90786.78	8997.16	42100.99	27531.43
C4-Fluoranthenes/Pyrenes	43675.06	5050.3	22284.19	27331.43 U
C0-Benzo(b)naphthothiophenes	95761.46	4337.03	625856.11	575552.01
C1-Benzo(b)naphthothiophenes	39637.36	4150.54	89078.51	68284.05
C2-Benzo(b)naphthothiophenes	12526.87	2683.1	24192.79	14774.05
C3-Benzo(b)naphthothiophenes	12395.15	1688.02	54788.21	47679.14
C4-Benzo(b)naphthothiophenes	2104.03	465.72	3861.4	2730.45
Benzo(a)anthracene	1354796.64	19761.41	2814695.87 D	2782388.99
Chrysene	1292269.51	28241.94	4315321.21 D	3838732.39 D

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Client ID	GWP07T13	GWP07T14	GWP07S01	GWP07S02
Battelle ID	R5536-P	R5537-P	R5538-P	R5539-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/12/07	01/12/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/09/07	02/09/07	02/09/07	02/08/07
Analytical Instrument	MS	MS	MS	MS
% Moisture	12.18	7.01	13.85	10.23
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.83	1.95	1.75	1.80
Size Unit-Basis	G_DRY	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit	384.24	12.91	1004.51	1030.86
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
C1-Chrysenes	350632.98	14801.48	334128.56	259854.13
C2-Chrysenes	115498.52	8152.3	82098.52	59322.94
C3-Chrysenes	84453.72	3756.19	19307.78	12665.37
C4-Chrysenes	15108.51	1094.7	U	12003.37 U
Benzo(b)fluoranthene	1046794.12	23921.92	3939621.07 D	4198747.8 D
Benzo(k)fluoranthene	1249814.43	24395.92	3111284.69	3214063.68
Benzo(e)pyrene	1018539.66	27937.45	4949803.45 D	5010843.9 D
Benzo(a)pyrene	1350021.51 D	35234.46	7607195.93 D	8192228.11 D
Pervlene	588546.04	10279.86	1960653.87 D	3213617.69
Indeno(1,2,3-cd)pyrene	992432.6 D	35802.93	7385669.2 D	7867281.06 D
Dibenz(a,h)anthracene	224314.97	5396.98	715519.56	599913.8
Benzo(g,h,i)perylene	1200172.25 D	38276.68 D	11127333.87 D	10804525.44 D
Total PAH	63373822.23	514633.54	170317016.4	143901804.8
Surrogate Recoveries (%)				
Naphthalene-d8	97	66	107	99
Acenaphthene-d10	62	76	66	63
Phenanthrene-d10	89	80	80	91
Benzo(a)pyrene-d12	181 NME	83	155 NME	139 NME

The Business of Innovation

Client ID	GWP07S03	GWP07S04	TDW3-4.5
Battelle ID	R5540-P	R5541-P	R5542-P
Sample Type	SA	SA	SA
Collection Date	01/12/07	01/12/07	09/26/06
Extraction Date	01/24/07	01/24/07	01/24/07
Analysis Date	02/12/07	02/08/07	02/09/07
Analytical Instrument	MS	MS	MS
% Moisture	20.14	12.16	14.86
% Lipid	NA	NA	NA
Matrix	TAR	TAR	SOIL
Sample Size	1.66	1.85	17.17
Size Unit-Basis	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit	1117.8	1337.34	6.48
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY
C3-Alkylbenzenes	1294243.87	158106.44 J	17.7 JT
C4-Alkylbenzenes	797410.21	382114.16	18.1 JT
C5-Alkylbenzenes	121385.55 J	67650.97 J	9.96 JT
C6-Alkylbenzenes	41402.02 J	33688.64 J	9.52 JT
Benzo(b)thiophene	1788561.97	2254325.92	328.76 T
C1-benzo(b)thiophenes	2045245.44	3058073.54	100.13 T
C2-benzo(b)thiophenes	1583925.71	2374427.64	102.97 T
C3-benzo(b)thiophenes	579130.31	763348.46	77 T
C4-benzo(b)thiophenes	116971.38	131691.56	40.58 T
Naphthalene	31350491.1 D	28051647.36 D	2965.03 T
C1-Naphthalenes	15885043.89	18087022.6	756.44 T
C2-Naphthalenes	8946938.73	9894089.64	642.61 T
C3-Naphthalenes	3262299.36	3138374.42	515.15 T
C4-Naphthalenes	723383.94	594581.58	332.96 T
Biphenyl	1314448.7	1419183.18	709.08 T
C1-Biphenyls + Dibenzofuran	1068258.12	990601.92	332.42 T
C2-Biphenyls + C1-Dibenzofurans	1131285.97	913603.05	295.76 T
Acenaphthylene	4151973.75 D	3725782.99	2665.3 T
Acenaphthene	355568.86	308642.76	113 T
Dibenzofuran	327097.45	299699.54	205.07 T
Fluorene	2469334.06	2089016.85	456.79 T
C1-Fluorenes	1981109.01	1635741.32	280.93 T
C2-Fluorenes C3-Fluorenes	1008404.37 335631.47	833572.19	365.9 T 1220.15 T
C4-Fluorenes	147321.1	439976.77 253597.8	572.27 T
Anthracene	2575820.2	1878925.68	3596.5 T
Phenanthrene	8251686.47 D	8665001.32 D	10905.75 T
C1-Phenanthrenes/Anthracenes	6530972.32	6966200.73	5353.75 T
C2-Phenanthrenes/Anthracenes	2876421.64	3716568.09	3710.04 T
C3-Phenanthrenes/Anthracenes	784697.31	1228739.4	1707.85 T
C4-Phenanthrenes/Anthracenes	170962.04	260004.02	530.66 T
Retene	U	U	490.91 T
Dibenzothiophene	1012181.57	1533863.59	1711.31 T
C1-Dibenzothiophenes	1200051.65	2093835.6	1166.67 T
C2-Dibenzothiophenes	753681.47	1646871.81	1188.44 T
C3-Dibenzothiophenes	301581.59	732821.53	681.43 T
C4-Dibenzothiophenes	68117.41	165057.12	196.34 T
Fluoranthene	1873682.81	1856741.01	48371.32 DT
Pyrene	3791787.81	3955992.67	69765.27 DT
C1-Fluoranthenes/Pyrenes	4310483.13	4455074.97	11962.25 T
C2-Fluoranthenes/Pyrenes	1666681.85	2144404.39	5261.06 T
C3-Fluoranthenes/Pyrenes	570324.57	778142.29	1646.56 T
C4-Fluoranthenes/Pyrenes	115630.39	166123.68	812.45 T
C0-Benzo(b)naphthothiophenes	234913.79	490166.25	5655.43 T
C1-Benzo(b)naphthothiophenes	445315.32	994243.91	2287.75 T
C2-Benzo(b)naphthothiophenes	236901.37	638001.83	940.77 T
C3-Benzo(b)naphthothiophenes	101327.36	248438.76	705.23 T
C4-Benzo(b)naphthothiophenes	27841.45	63057.34	126.43 T
Benzo(a)anthracene	1484981.4	1613574.99	17677.03 T
Chrysene	1614197.23	1754172.46	21363.2 T

Client ID	GWP07S03	GWP07S04	TDW3-4.5
Battelle ID	R5540-P	R5541-P	R5542-P
Sample Type	SA	SA	SA
Collection Date	01/12/07	01/12/07	09/26/06
Extraction Date	01/24/07	01/24/07	01/24/07
Analysis Date	02/12/07	02/08/07	02/09/07
Analytical Instrument	MS	MS	MS
% Moisture	20.14	12.16	14.86
% Lipid	NA	NA	NA
Matrix	TAR	TAR	SOIL
Sample Size	1.66	1.85	17.17
Size Unit-Basis	G DRY	G DRY	G DRY
Minimum Reporting Limit	1117.8	1337.34	6.48
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY
C1-Chrysenes	1650064.43	2194426.25	5462.79 T
C2-Chrysenes	699503.7	1208945.48	2180.9 T
C3-Chrysenes	314904.32	421187.55	793.81 T
C4-Chrysenes	68838.45	91294.52	735.01 T UT
Benzo(b)fluoranthene	520198.7	567719.71	21099.37 T
Benzo(k)fluoranthene	759851.51	793548.85	19967.55 T
Benzo(e)pyrene	746671.21	1082489.22	23771.27 T
Benzo(a)pyrene	1448281.84	1507281.33	24775.08 DT
Perylene	262250.29	298368.92	8679.4 T
Indeno(1,2,3-cd)pyrene	522843.71	590739.46	23701.94 DT
Dibenz(a,h)anthracene	203297.51	224563.2	4440.35 T
Benzo(g,h,i)perylene	545106.51	808491.22	28198.79 DT
Total PAH	119691849.3	125752378.6	381095.09 T
Surrogate Recoveries (%)			
Naphthalene-d8	77	103	78
Acenaphthene-d10	74	63	87
Phenanthrene-d10	77	82	86
Benzo(a)pyrene-d12	219 NME	169 NME	101

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Client ID	Procedural Blank	
Battelle ID	BJ939PB-P	
Sample Type	PB	
Collection Date	01/24/07	
Extraction Date	01/24/07	
Analysis Date	02/10/07	
Analytical Instrument	MS	
% Moisture	19.52	
% Lipid	NA	
Matrix	SOIL, TAR	
Sample Size	2.44	
Size Unit-Basis	G_DRY	
Minimum Reporting Limit	8.21	
Units	NG/G_DRY	
	o (o)	
C3-Alkylbenzenes	3.48 J	
C4-Alkylbenzenes	U	
C5-Alkylbenzenes	U	
C6-Alkylbenzenes	U	
Benzo(b)thiophene	U	
C1-benzo(b)thiophenes	U	
C2-benzo(b)thiophenes	U	
C3-benzo(b)thiophenes	U	
C4-benzo(b)thiophenes	U 8.42 N	
Naphthalene	8.42 N 2.13 J	
C1-Naphthalenes	2.13 J U	
C2-Naphthalenes	UU	
C3-Naphthalenes	UU	
C4-Naphthalenes Biphenyl	U	
C1-Biphenyls + Dibenzofuran	U	
C1-Diphenyls + C1-Dibenzofurans	U	
Acenaphthylene	U	
Acenaphthene	U	
Dibenzofuran	U	
Fluorene	U	
C1-Fluorenes	Ű	
C2-Fluorenes	Ŭ	
C3-Fluorenes	Ŭ	
C4-Fluorenes	Ŭ	
Anthracene	U	
Phenanthrene	2.37 J	
C1-Phenanthrenes/Anthracenes	U	
C2-Phenanthrenes/Anthracenes	Ū	
C3-Phenanthrenes/Anthracenes	U	
C4-Phenanthrenes/Anthracenes	U	
Retene	U	
Dibenzothiophene	U	
C1-Dibenzothiophenes	U	
C2-Dibenzothiophenes	U	
C3-Dibenzothiophenes	U	
C4-Dibenzothiophenes	U	
Fluoranthene	1.34 J	
Pyrene	1.54 J	
C1-Fluoranthenes/Pyrenes	U	
C2-Fluoranthenes/Pyrenes	U	
C3-Fluoranthenes/Pyrenes	U	
C4-Fluoranthenes/Pyrenes	U	
C0-Benzo(b)naphthothiophenes	U	
C1-Benzo(b)naphthothiophenes	U	
C2-Benzo(b)naphthothiophenes	U	
C3-Benzo(b)naphthothiophenes	U	
C4-Benzo(b)naphthothiophenes	U	
Benzo(a)anthracene	U	
Chrysene	U	

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Client ID	Procedural Blank	
Battelle ID	BJ939PB-P	
Sample Type	PB	
Collection Date	01/24/07	
Extraction Date	01/24/07	
Analysis Date	02/10/07	
Analytical Instrument	MS	
% Moisture	19.52	
% Lipid	NA	
Matrix	SOIL, TAR	
Sample Size	2.44	
Size Unit-Basis	G_DRY	
Minimum Reporting Limit	8.21	
Units	NG/G_DRY	
C1-Chrysenes	U	
C2-Chrysenes	U	
C3-Chrysenes	U	
C4-Chrysenes	U	
Benzo(b)fluoranthene	U	
Benzo(k)fluoranthene	U	
Benzo(e)pyrene	U	
Benzo(a)pyrene	U	
Perylene	U	
Indeno(1,2,3-cd)pyrene	1.51 J	
Dibenz(a,h)anthracene	0.73 J	
Benzo(g,h,i)perylene	2.07 J	
Total PAH	20.11 J	
Surrogate Recoveries (%)		
Naphthalene-d8	94	
Acenaphthene-d10	89	
Phenanthrene-d10	94	
Benzo(a)pyrene-d12	54 71	
Donzo(a)pyrene-u rz	71	

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	060208-03: Sand,			
Client ID	White Quartz, -50+70			
Battelle ID	BJ940LCS-P			
Sample Type	LCS			
Collection Date	01/24/07			
Extraction Date	01/24/07			
Analysis Date	02/10/07			
Analytical Instrument	MS			
% Moisture	NA			
% Lipid	NA			
Matrix	SOIL, TAR			
Sample Size	20.01			
Size Unit-Basis	G_DRY			
Minimum Reporting Limit	1.00			
Units	NG/G_DRY	Target %	Recovery Qualifier	
01113	NO/O_DIT	Target /d	Recovery Qualifier	
C3-Alkylbenzenes	U			
C4-Alkylbenzenes	U			
C5-Alkylbenzenes	U			
C6-Alkylbenzenes	U			
Benzo(b)thiophene	132.09	125.02	106	
C1-benzo(b)thiophenes	U			
C2-benzo(b)thiophenes	U			
C3-benzo(b)thiophenes	U			
C4-benzo(b)thiophenes	U			
Naphthalene	138.71	124.96	111	
C1-Naphthalenes	U			
C2-Naphthalenes	U			
C3-Naphthalenes	Ŭ			
C4-Naphthalenes	Ŭ			
Biphenyl	134.58	125.15	108	
C1-Biphenyls + Dibenzofuran	U	120.10	100	
C2-Biphenyls + C1-Dibenzofurans	Ŭ			
Acenaphthylene	129.1	125.05	103	
Acenaphthene	136.48	125.02	109	
Dibenzofuran	135.53	125.17	109	
Fluorene	137.31	125.01	110	
C1-Fluorenes	137.31 U	125.01	110	
C2-Fluorenes	U			
C3-Fluorenes	U			
C4-Fluorenes	U			
Anthracene	134.3	124.96	107	
Phenanthrene	143.66		115	
C1-Phenanthrenes/Anthracenes	143.00 U	125.00	115	
C2-Phenanthrenes/Anthracenes	U			
C3-Phenanthrenes/Anthracenes	U			
C4-Phenanthrenes/Anthracenes	U			
Retene	U	405 54	440	
Dibenzothiophene	137.72	125.51	110	
C1-Dibenzothiophenes	U			
C2-Dibenzothiophenes	U			
C3-Dibenzothiophenes	U			
C4-Dibenzothiophenes	U			
Fluoranthene	145.74	125.00	117	
Pyrene	142.26	124.98	114	
C1-Fluoranthenes/Pyrenes	U			
C2-Fluoranthenes/Pyrenes	U			
C3-Fluoranthenes/Pyrenes	U			
C4-Fluoranthenes/Pyrenes	U			
C0-Benzo(b)naphthothiophenes	U			
C1-Benzo(b)naphthothiophenes	U			
C2-Benzo(b)naphthothiophenes	U			
C3-Benzo(b)naphthothiophenes	U			
C4-Benzo(b)naphthothiophenes	U	(a ·		
Benzo(a)anthracene	108.23	124.97	87	

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	060208-03: Sand,				
Client ID	White Quartz, -50+70				
Battelle ID	BJ940LCS-P				
Sample Type	LCS				
Collection Date	01/24/07				
Extraction Date	01/24/07				
Analysis Date	02/10/07				
Analytical Instrument	MS				
% Moisture	NA				
% Lipid	NA				
Matrix	SOIL, TAR				
Sample Size	20.01				
Size Unit-Basis	G_DRY				
Minimum Reporting Limit	1.00				
Units	NG/G_DRY		Target %	Recovery	Qualifier
Chrysene	119.18		124.99	95	
C1-Chrysenes		U			
C2-Chrysenes		U			
C3-Chrysenes		U			
C4-Chrysenes		U			
Benzo(b)fluoranthene	107.83		125.04	86	
Benzo(k)fluoranthene	121.31		125.01	97	
Benzo(e)pyrene	114.48		125.25	91	
Benzo(a)pyrene	105.11		125.04	84	
Perylene	88.5		125.19	71	
Indeno(1,2,3-cd)pyrene	101.19		125.00	81	
Dibenz(a,h)anthracene	109.51		125.01	88	
Benzo(g,h,i)perylene	110.44		124.98	88	
Total PAH	2601.17				
Surrogate Recoveries (%)					
Nanhthalana da	100				

Naphthalene-d8	109
Acenaphthene-d10	103
Phenanthrene-d10	112
Benzo(a)pyrene-d12	76

The Business of Innovation

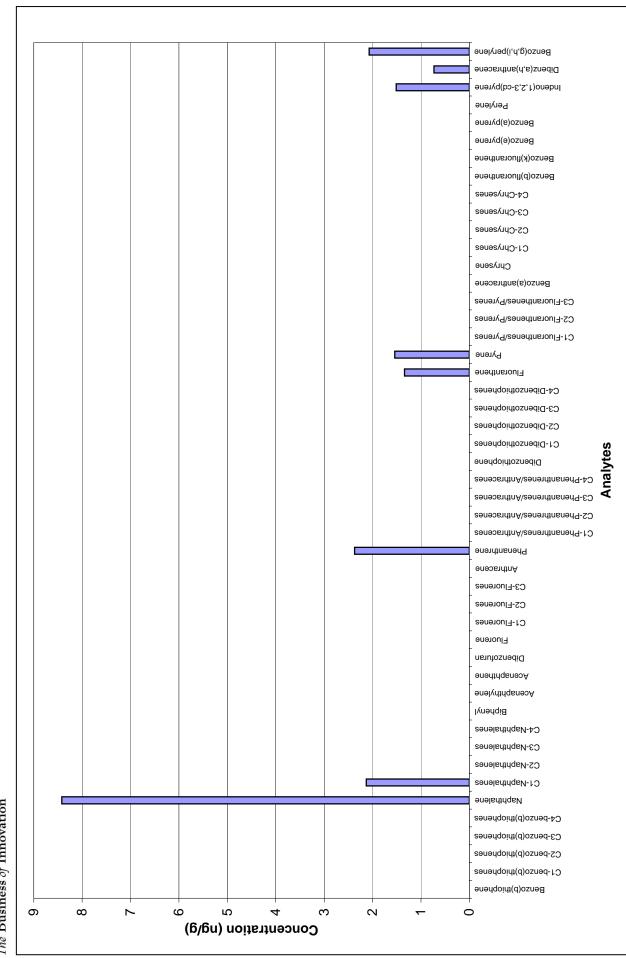
	ONICO: North Olars				
Client ID	GN62: North Slope Crude				
Client ID	Ciude				
Battelle ID	BJ959NSC-P				
Sample Type	NSC				
Collection Date	01/30/07				
Extraction Date	01/30/07				
Analysis Date	02/07/07				
Analytical Instrument	MS				
% Moisture	NA				
% Lipid	NA				
Matrix	OIL				
Sample Size	5.01				
Size Unit-Basis	G_OIL				
Minimum Reporting Limit	1.3				
Units	NG/G_OIL		Target % I	Difference	Qualifier
··· ··					
C3-Alkylbenzenes	2027.44				
C4-Alkylbenzenes	1510.05				
C5-Alkylbenzenes	795.79				
C6-Alkylbenzenes	488.99				
Benzo(b)thiophene	13.52				
C1-benzo(b)thiophenes	45				
C2-benzo(b)thiophenes	79.93		95.74	16.5	
C3-benzo(b)thiophenes	141.48		132.67	6.6	
C4-benzo(b)thiophenes	96.21		96.72	0.5	
Naphthalene	806.17		740.29	8.9	
C1-Naphthalenes	1634.24		1516.04	7.8	
C2-Naphthalenes	2010.9		2000.10	0.5	
C3-Naphthalenes	1445.89		1526.96	5.3	
C4-Naphthalenes	785.13		898.03	12.6	
Biphenyl	371.59		220.02	0.0	
C1-Biphenyls + Dibenzofuran	240.71		220.82	9.0	
C2-Biphenyls + C1-Dibenzofurans	514.5	U			
Acenaphthylene Acenaphthene	13.75	0	14.50	5.2	
Dibenzofuran	77.57		77.75	0.2	
Fluorene	96.04		92.51	3.8	
C1-Fluorenes	223.14		227.01	3.8 1.7	
C2-Fluorenes	345.32		367.09	5.9	
C3-Fluorenes	295.4		326.32	9.5	
C4-Fluorenes	212.42		020.02	0.0	
Anthracene	212.12	U			
Phenanthrene	285.03	Ũ	249.49	14.2	
C1-Phenanthrenes/Anthracenes	584.34		549.17	6.4	
C2-Phenanthrenes/Anthracenes	669.09		642.72	4.1	
C3-Phenanthrenes/Anthracenes	453.02		446.11	1.5	
C4-Phenanthrenes/Anthracenes	176.09		180.02	2.2	
Retene	74.51				
Dibenzothiophene	241.56		210.35	14.8	
C1-Dibenzothiophenes	440.7		409.03	7.7	
C2-Dibenzothiophenes	573.69		551.46	4.0	
C3-Dibenzothiophenes	483.6		471.36	2.6	
C4-Dibenzothiophenes	261.9		243.11	7.7	
Fluoranthene		U			
Pyrene	14.53		12.99	11.9	
C1-Fluoranthenes/Pyrenes	84.98		70.92	19.8	
C2-Fluoranthenes/Pyrenes	145.97		117.89	23.8	
C3-Fluoranthenes/Pyrenes	154.91		137.25	12.9	
C4-Fluoranthenes/Pyrenes	123.35				
C0-Benzo(b)naphthothiophenes	46.42				
C1-Benzo(b)naphthothiophenes	162.78				
C2-Benzo(b)naphthothiophenes	204.91				
C3-Benzo(b)naphthothiophenes	163.24				
C4-Benzo(b)naphthothiophenes	66.74				
Benzo(a)anthracene		U			

The Business of Innovation

	GN62: North Slope			
Client ID	Crude			
Battelle ID	BJ959NSC-P			
Sample Type	NSC			
Collection Date	01/30/07			
Extraction Date	01/30/07			
Analysis Date	02/07/07			
Analytical Instrument	MS			
% Moisture	NA			
% Lipid	NA			
Matrix	OIL			
Sample Size	5.01			
Size Unit-Basis	G_OIL			
Minimum Reporting Limit	1.3			
Units	NG/G_OIL	Target % D	ifference	Qualifier
Chrysene	51.12	47.18	8.4	
C1-Chrysenes	85.28	78.82	8.2	
C2-Chrysenes	113.52	102.67	10.6	
C3-Chrysenes	97.23	85.36	13.9	
C4-Chrysenes	62.06	61.99	0.1	
Benzo(b)fluoranthene	6.18	6.08	1.6	
Benzo(k)fluoranthene	U			
Benzo(e)pyrene	13.86	12.88	7.6	
Benzo(a)pyrene	U			
Perylene	U			
Indeno(1,2,3-cd)pyrene	U			
Dibenz(a,h)anthracene	1.1 J			
Benzo(g,h,i)perylene	3.86	3.44	12.2	
Total PAH	12977.88			
Surragata Pasavarias (%)				
Surrogate Recoveries (%)				

Naphthalene-d8	109
Acenaphthene-d10	106
Phenanthrene-d10	97
Benzo(a)pyrene-d12	119

Procedural Blank (BJ939PB-P)



GWP07T01 (R5521-P)

GWP07T02 (R5525-P)

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								C2-Fluorenes C3-Fluorenes C3-Fluorenes Phenanthrenes/Phinacenes C3-Phenanthrenes/Phinacenes C3-Phenanthrenes/Phinacenes C4-Phenanthrenes/Phinacenes C3-Diberzothiophenes C3-Diberzothiophenes C3-Diberzothiophenes C3-Diberzothiophenes	Analytes
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GWP07T03 (R5526-P)

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						Benzo(g,h,i)perylene	
						Dibenz(a,h)anthracene	
						Indeno(5,2,5)pyrene	
						- Perylene	
						Benzo(a)pyrene	
						Benzo(e)pyrene	
						Benzo(k)fluoranthene	
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						C4-Chrysenes	
						C3-CµLysenes	
						C2-Chrysenes	
						C1-Chrysenes	
						Chrysene	
						Benzo(a)anthracene	
						C3-Fluoranthenes/Pyrenes	
						C2-Fluoranthenes/Pyrenes	
						C1-Fluoranthenes/Pyrenes	
						Pyrene	
						Fluoranthene	
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						C3-Dibenzothiophenes	
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						C1-Dibenzothiophenes	nalytes
						Dibenzothiophene	ž
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						C2-Phenanthrenes/Anthracenes	
						Phenanthrenes/Anthracenes	
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						C1-Fluorenes	
						Fluorene	
						Dibenzofuran	
						- -	
						Acenaphthylene	
						Biphenyl	
						C4-Naphthalenes	
						C3-Naphthalenes	
						C2-Naphthalenes	
						C1-Naphthalenes	
						Asphthalane Vaphthalene	
						C4-benzo(b)thiophenes	
						C3-penzo(b)thiophene	
						C2-benzo(b)thiophene c	
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						Benzo(b)thiophene	
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GWP07T04 (R5527-P)

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						ا _	- C2-Naphthalenes	
							C3-Naphthalenes	
							C4-Naphthalenes	
						I	Biphenyl	
							Acenaphthylene	
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							Dibenzofuran	
							Fluorene	
							C1-Fluorenes	
							C2-Fluorenes	
							C3-Fluorenes	
							Anthracene	
							Phenanthrene	
						ſ	canenativenes/Anthracenes	
							cenes/Anthracenes/Anthracenes	
							canenathrenes/Anthracenes	A
							C4-Phenanthrenes/Anthracenes	Analytes
							Dibenzothiophene	yte
							C-1-Dibenzothiophenes	ŝ
							C2-Dibenzothiophenes	
							C3-Dibenzothiophenes	
							C4-Dibenzothiophenes	
							Fluoranthene	
							- - -	
						г	C1-Fluoranthenes/Pyrenes	
							C2-Fluoranthenes/Pyrenes	
						L	C3-Fluoranthenes/Pyrenes	
						L T	Benzo(a)anthracene	
							C1-Chrysene	
							C2-Chrysenes	
							C3-Chrysenes	
							C4-Chrysenes	
						I	Benzo(b)fluoranthene Benzo(b)fluoranthene	
						L r	-	
						I T	Benzo(k)fluoranthene - -	
							Benzo(e)oznaB	
							Bendenzo(a)pyrene 	
							Perylene - -	
							Indeno(1,2,3-cd)pyrene	
							Dibenz(a,h)anthracene	

GWP07T05 (R5528-P)

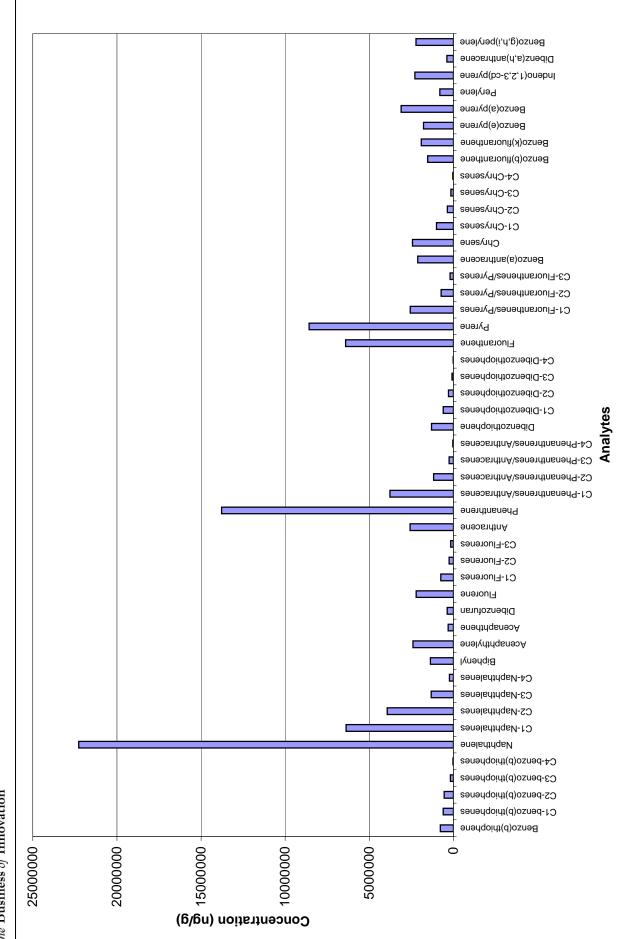
GWP07T06 (R5529-P)

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					C3-Phenanthrenes/Anthracenes C2-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes
					Pyrene Pyrenes Fororanthenes/Pyrenes C2-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes Chrysenes Benzo(b)fluoranthene Perylene

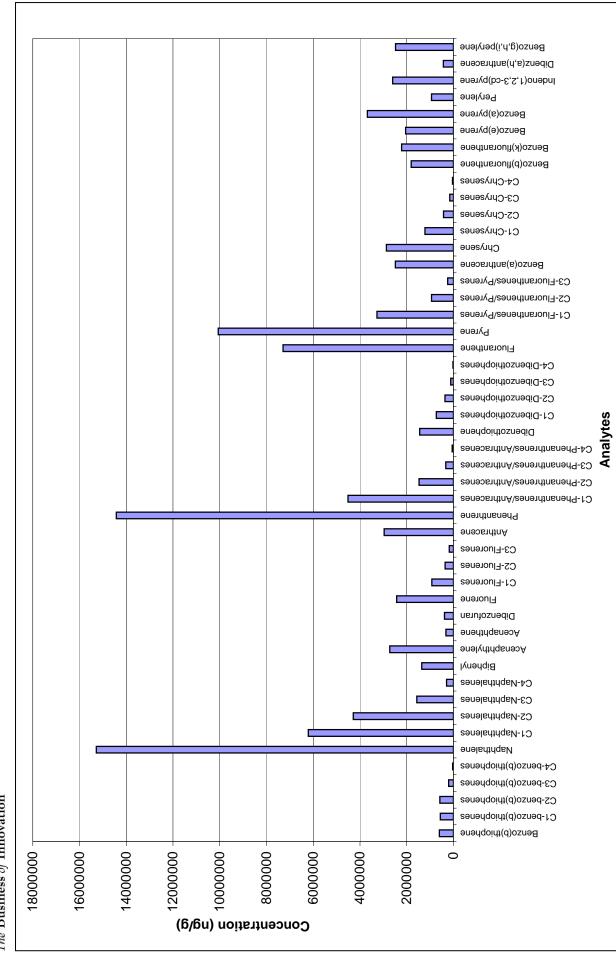
GWP07T07 (R5530-P)

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C2-Photenes/Prenes/ C3-Photomhenes/ C3-Photomhenes/ C3-Photomhenes/ C3-Photomhenes/Prenes/ C3-Photomhenes/Prenes/ C3-Photomhenes/Prenes/ C3-Photomhenes/Prenes/ C3-Photomhenes/Prenes/ C3-Photomhenes/Prenes/ C3-Photomhenes/Prenes/ C3-Photomhenes/Prenes/ C3-Photomhenes/Prenes/ C3-Photomhenes/Prenes/ C3-Photomhenes/Prenes/ C3-Photomhenes/Prenes/ C3-Photomhenes/ C3-Photomhenes/ C3-Photomhenes/ C3-Photomhenes/ C3-Photomh										-
C2-Fluorente C3-Fluorente C3-Fluorente C3-Fluorente C3-Fluorente C3-Fluorente C3-Fluorente C3-Fluorente C3-Chrysen										-
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C2-Diberothiophenes C3-Tiboranthenes/Pyrenes C3-Tiboranthenes/Pyrenes C3-Tiboranthenes/Pyrenes C3-Chrysenes Elenzo(a)pyrene Perviene Diberz(a,h)anthracen									1	C3-Phenanthrenes/Anthracenes
C2-Diberothiophenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes Elenzo(a)pyrene Perylene Pery										C4-Phenanthrenes/Anthracenes
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Pyrene C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Chrysenes C3-C										-
C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Chrysenes C4-Chrysenes C4-Chrysenes Benzo(s)pyrene Perylene Perylene Dibenz(s,h)anthracene										-
C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthene C4-Chrysenes C3-Fluoranthene C4-Chrysenes C4-Chr										-
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C3-Chrysenes C3-Chrysenes Benzo(b)fluoranthene Benzo(b)fluoranthene Benzo(k)fluoranthene Benz										-
C4-Chrysenes Benzo(k)fluoranthene Benzo(k)pyrene Perylene Dibenz(a,h)anthracene										-
Benzo(b)tronomic (a, h) substraint (a, h) substraint (a) (b) constraint (b) (b) constraint (c)										-
Benzo(k)fluoranthrene Benzo(k)pyrene Perylene Perylene Dibenz(a,h)tustication Dibenz(a,h)tu										-
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Diberof(a,h)anthracene										Benzo(a)pyrene
Dibenzich, h) anthracene										Leuylene
										Indeno(1,2,3-cd)pyrene
geuzo(â'u'ı)beu\leue										-
										Benzo(g,h,i)perylene

GWP07T08 (R5531-P)

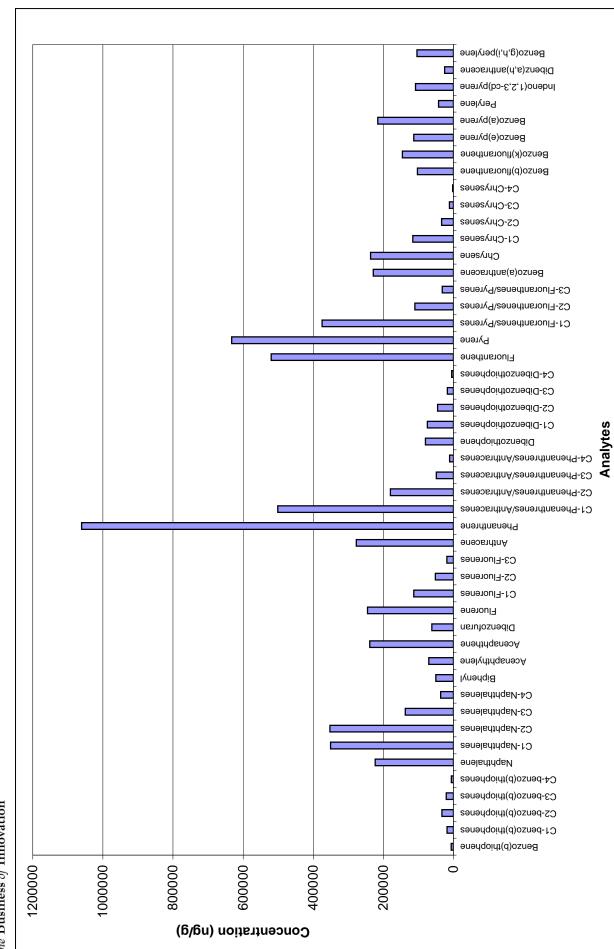


GWP07T09 (R5532-P)



GWP07T10 (R5533-P)

GWP07T11 (R5534-P)



GWP07T12 (R5535-P)

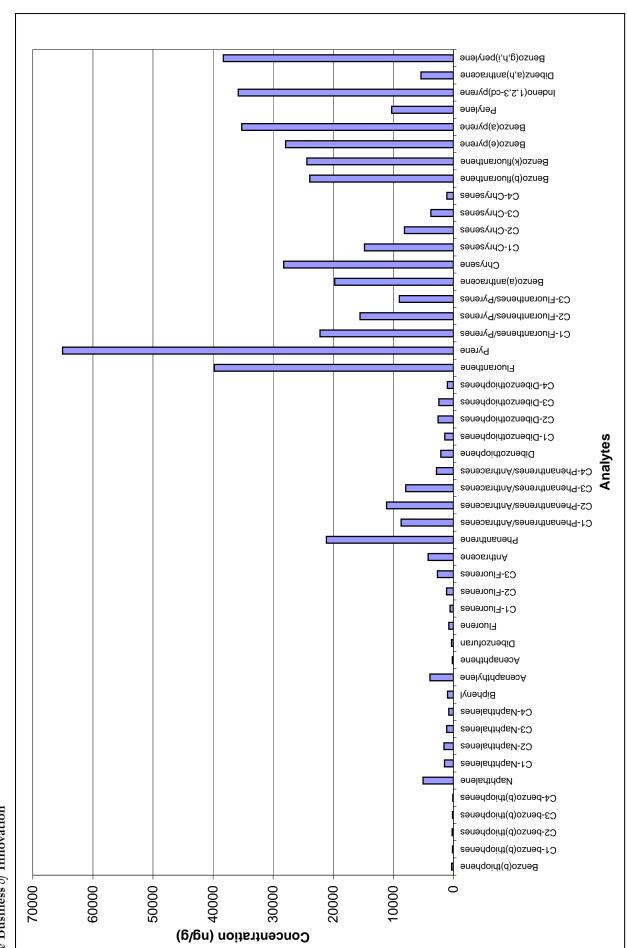
					Benzo(e)pyrene Benzo(a)pyrene Perylene Perylene Prene Dibenz(a,h)anthracene
					Benzo(k)fluoranthene Benzo(b)fluoranthene Benzo(k)fluoranthene
					C3-Fluoranthenes/Pyrenes Benzo(a)anthracene Chrysenes C1-Chrysenes
					C2-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes
					C3-Phenanthrenes/Anthracenes C4-Phenanthrenes/Anthracenes C1-Dibenzothiophenes C2-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes
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GWP07T13 (R5536-P)

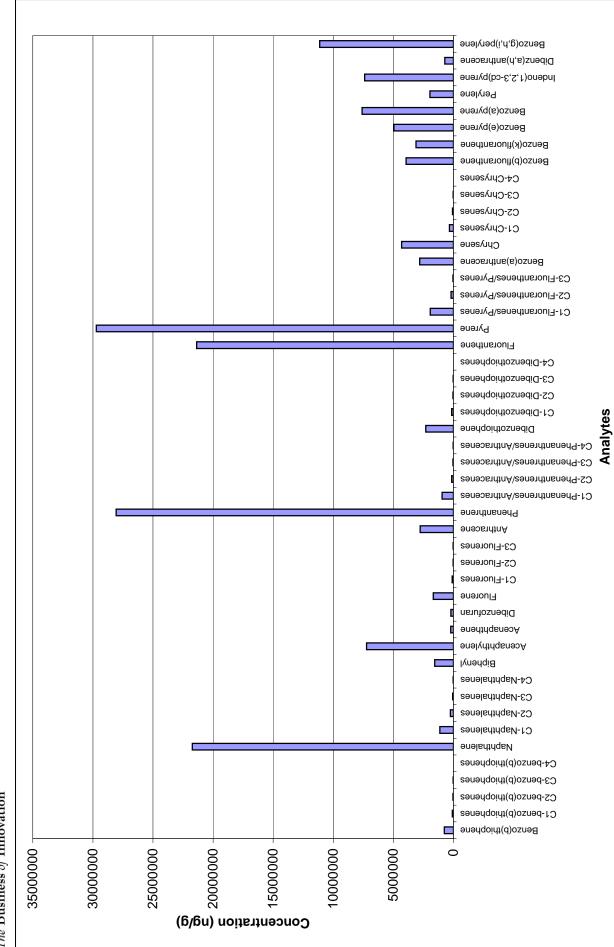
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								Fluoranthene Pyrenes C1-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes Benzo(a)anthracene
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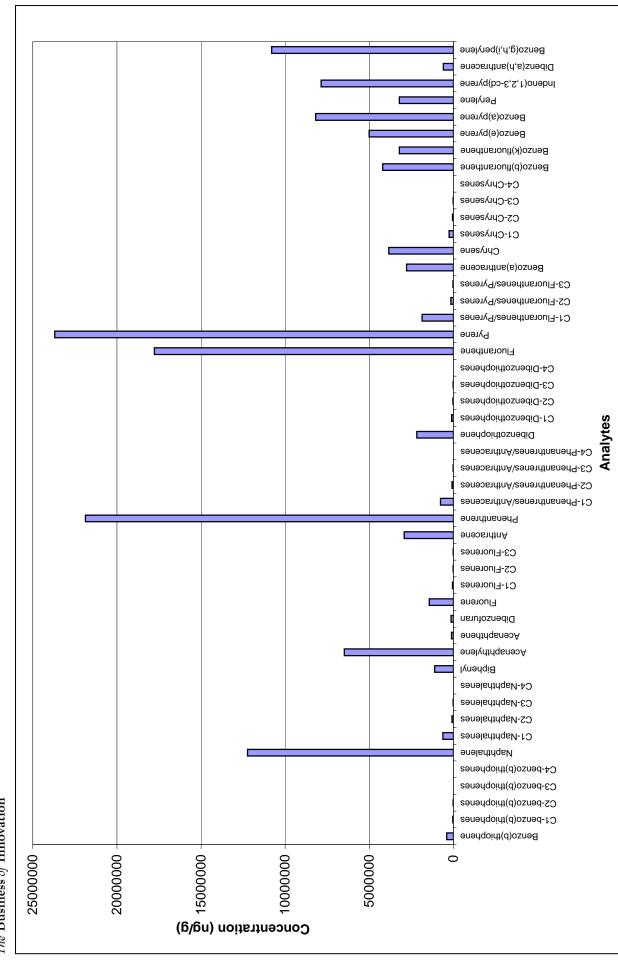
GWP07T14 (R5537-P)



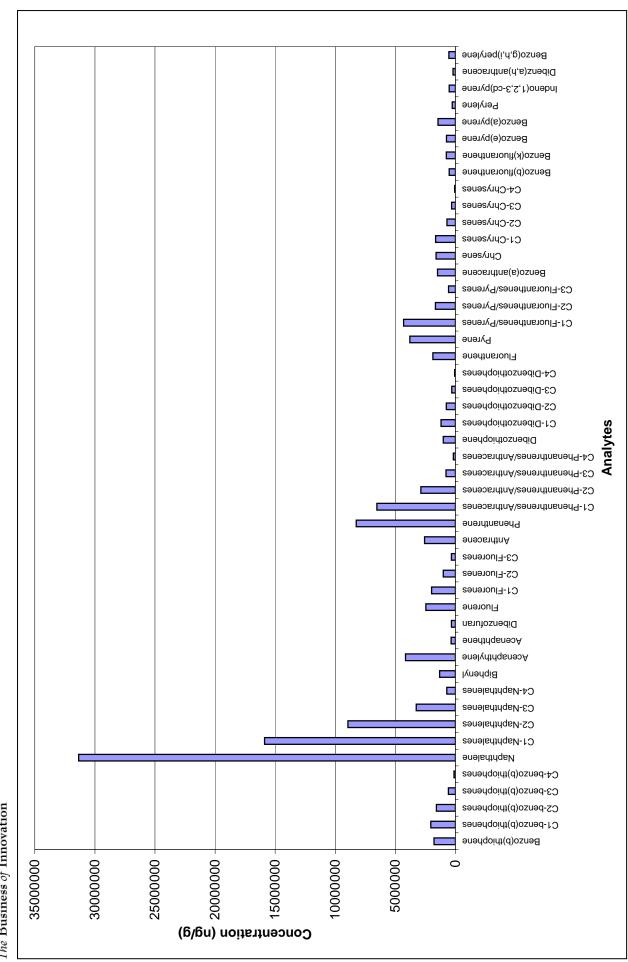
GWP07S01 (R5538-P)



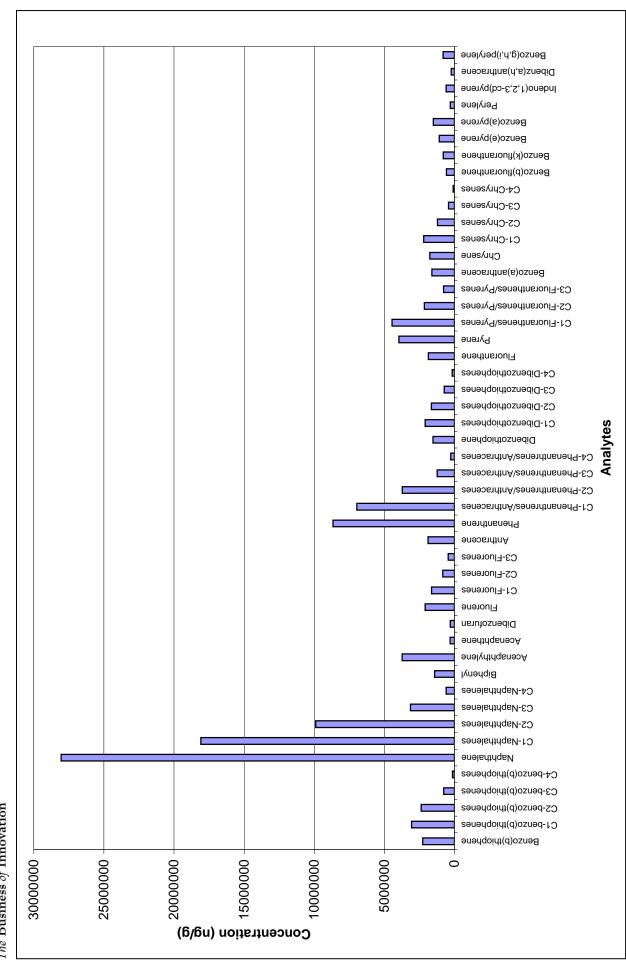
GWP07S02 (R5539-P)



GWP07S03 (R5540-P)



GWP07S04 (R5541-P)

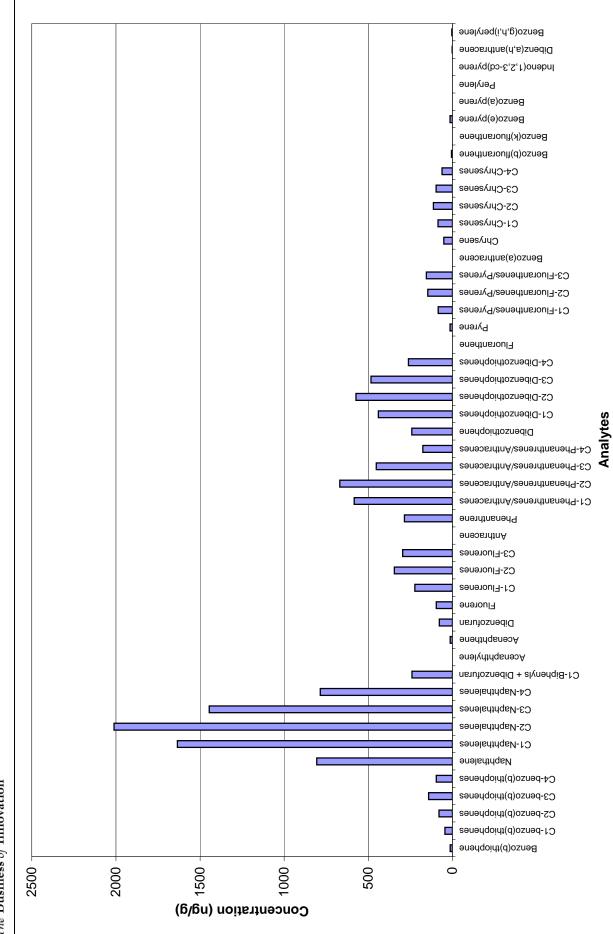


TDW3-4.5 (R5542-P)

80000					Benzo(b)thiophenee C1-benzo(b)thiophenes C2-benzo(b)thiophenes C3-benzo(b)thiophenes C4-benzo(b)thiophenes C1-Naphthalenes C1-Naphthalenes C3-Naphthalenes C3-Naphthalenes
					Benzo(b)thiophenes C1-benzo(b)thiophenes C2-benzo(b)thiophenes C3-benzo(b)thiophenes C4-benzo(b)thiophenes C4-benzo(b)thiophenes C1-Naphthalenes C3-Naphthalenes C3-Naphthalenes Biphanyl Biphanyl C4-Ribhene Fluorene C1-Fluorenes C1-Fluorenes
					C1-benzo(b)thiophenes C2-benzo(b)thiophenes C3-benzo(b)thiophenes C3-benzo(b)thiophenes C4-benzo(b)thialenes C2-Vaphthalenes C3-Vaphthalenes C3-Vaphthalenes C3-Vaphthalenes Biphenyl Biphenyl Biphenyl C4-fluorene C1-Fluorenes C1-Fluorenes
					C2-benzo(b)thiophenes C3-benzo(b)thiophenes C4-benzo(b)thiophenes C4-benzo(b)thiophenes C4-benzo(b)thialenes C3-Naphthalenes C4-Naphthalenes C4-Naphthalenes Biphenyl Biphenyl C4-Fluorene C1-Fluorenes C3-Fluorenes
					C3-benzo(b)thiophenes C4-benzo(b)thiophenes V4sphthalenes C1-V4sphthalenes C2-V4sphthalenes C3-V4sphthalenes Biphenyl Biphenyl Acenaphthylene Pibenzofuran Fluorene C1-Fluorenes C3-Fluorenes
					C4-benzo(b)thiophenes Maphthalenes C3-Naphthalenes C3-Naphthalenes C3-Naphthalenes C4-Naphthalenes Biphenyl Biphenyl Acenaphthylene C4-Fluorene C1-Fluorenes C3-Fluorenes
					Naphthalene C1-Naphthalenes C2-Naphthalenes C3-Naphthalenes C4-Naphthalenes Biphenyl Acenaphthylene Dibenzofuran Fluorene C1-Fluorenes
					C1-Naphthalenes C2-Naphthalenes C3-Naphthalenes Biphenyl Acenaphthylene Dibenzofuran Fluorene C1-Fluorenes C1-Fluorenes
					C2-Naphihalenes C3-Naphihalenes C4-Naphihalenes Biphenyl Acenaphihylene Dibenzofuran Fluorene C1-Fluorenes
					C3-Naphihalenes C4-Naphihalenes Biphenyl Acenaphihylene Dibenzofuran Fluorene C1-Fluorenes C2-Fluorenes
					Biphenyl Biphenyl Acenaphthylene Acenaphthene Dibenzofuran Fluorene C1-Fluorenes C3-Fluorenes
					Acenaphthylene Acenaphthene Dibenzofuran Fluorene C1-Fluorenes C2-Fluorenes
					Acenaphthene Dibenzofuran Fluorene C1-Fluorenes C2-Fluorenes
					Dibenzofuran Fluorenes C1-Fluorenes
					Fluorenes C1-Fluorenes C2-Fluorenes
					C1-Fluorenes
					- C2-Fluorenes
		I			-
					C3-Fluorenes
		l			-
		l		-	- Anthracene
		l		Г	Phenanthrand Phenanthrand D
		l			C1-Phenanthrenes/Anthracenes
		l			C2-Phenanthrenes/Anthracenes
		l			C3-Phenanthrenes/Anthrecenes
		1			- Dipension of the contension
		1			Dibenzothornadi - SanandopidtoznadiC-tO
		1			C1-Dibenzothiophenes
		1			C2-Dibenzothiophenes
		l			C4-Dibenzothiophenes
					Fluoranthene
					- - -
		L			C1-Fluoranthenes/Pyrenes
		1			
		1			C3-Fluoranthenes/Pyrenes
		1			Benzo(a)anthracene
		1			Chrysene
		l			C1-Chrysenes
		1			C2-Chrysenes
		l			C3-Chrysenes
		l			C4-Chrysenes
		l		1	Benzo(b)fluoranthene
		l			Benzo(k)fluoranthene
		l			Benzo(e)pyrene
		1			Benzo(a)pyrene
		l			Perylene
		l			encectd*gc(d,c)zgodi
		l			Dibenz(a,h)anthracene - Benzo(g,h,i)perylene
					-



GN62: North Slope Crude (BJ959NSC-P)



The Business of Innovation

Client ID	GWP07T01	GWP07T02	GWP07T03	GWP07T04
Battelle ID	R5521-P	R5525-P	R5526-P	R5527-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/11/07	01/11/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/12/07	02/10/07	02/09/07	02/09/07
Analytical Instrument	MS	MS	MS	MS
% Moisture	28.88	50.1	50.3	34.93
% Lipid	NA	NA	NA	54.95 NA
Matrix	TAR	TAR	TAR	TAR
	1.44	1.02	1.01	1.44
Sample Size			G_DRY	
Size Unit-Basis	G_DRY	G_DRY		G_DRY
Minimum Reporting Limit	281.71	1242.82	1046.38	489.28
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
C23 Tricyclic Terpane	U	U	U	U
C29 Tricyclic Terpane -22S	U	U	U	U
C29 Tricyclic Terpane -22R	U	U	U	U
18a(H)-22,29,30-Trisnorneohopane -TS	U	U	U	U
	-	_	_	-
17a(H)-22,29,30-Trisnorhopane -TM	556.01 J	U	U	U
30-Norhopane	1062.22	U	U	U
18a(H) & 18b(H)-Oleananes	U	U	U	U
Hopane	1827.01	U	U	U
30-Homohopane -22S	U	U	U	U
30-Homohopane -22R	U	U	U	U
13b(H),17a(H)-20S-Diacholestane	U	U	U	U
13b(H),17a(H)-20R-Diacholestane	U	U	U	U
14a(H),17a(H)-20R-methylcholestane	519.25 J	U	U	478.26 J
14a(H),17a(H)-20S-Ethylcholestane	407.28 J	U	U	445.87 J
14a(H),17a(H)-20R-Ethylcholestane	362.99 J	U	U	316.3 J
C21-TAS	U	U	U	U
C26-TAS(20S)	81.59 J	213.02 J	U	U
C26,C27-TAS	205.25 J	432.58 J	257.21 J	U
C27-TAS(20R)	116.48 J	241.98 J	206.94 J	U
C28-TAS(20S)	82.36 J	174.71 J	U	U
C28-TAS(20R)	93.9 J	231.7 J	183.48 J	U
C21-MAS	U	U	U	U
C22-MAS	U	U	U	U
C27-MAS	U	U	U	U
C27-20R-MAS	3355.08	U	U	U
C27-20S-MAS	989.82	U	U	U
C28-20S-MAS	U	U	U	U
C27-C2920S/R-MAS	35605.15	U	Ŭ	U
C29-20S-MAS	44868.64	Ū	Ū	Ŭ
C29-20R-MAS	U	Ŭ	2221.04	Ŭ
TAS 245	NA	NA	NA	NA
MAS_239	NA	NA	NA	NA
-				

Surrogate Recoveries (%)									
5b(H)-Cholane	18404 NME	32515 NME	23623 NME	22622 NME					

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07T05	GWP07T06	GWP07T07	GWP07T08
Battelle ID	R5528-P	R5529-P	R5530-P	R5531-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/11/07	01/11/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
	02/08/07	02/08/07	02/08/07	01/24/07
Analysis Date				
Analytical Instrument	MS	MS	MS	MS
% Moisture	4.23	6.94	6.47	30.28
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.92	1.93	1.87	1.49
Size Unit-Basis	G_DRY	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit	1452.54	971.44	994.26	935.87
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
C23 Tricyclic Terpane	U	U	U	U
C29 Tricyclic Terpane -22S	U	U	U	U
C29 Tricyclic Terpane -22R	U	U	U	U
18a(H)-22,29,30-Trisnorneohopane -TS	U	U	U	U
17a(H)-22,29,30-Trisnorhopane -TM	U	U	U	U
30-Norhopane	5597.06	4469.68	4619.9	U
18a(H) & 18b(H)-Oleananes	U	U	U	Ŭ
Hopane	7476.45	5024.06	5375.22	U
30-Homohopane -22S	U	5024.00 U	3152.17	U
	U	U		U
30-Homohopane -22R			3078.5	_
13b(H),17a(H)-20S-Diacholestane	U	U	U	U
13b(H),17a(H)-20R-Diacholestane	U	U	U	U
14a(H),17a(H)-20R-methylcholestane	U	U	U	U
14a(H),17a(H)-20S-Ethylcholestane	U	U	1364.18 J	U
14a(H),17a(H)-20R-Ethylcholestane	U	U	1732.93 J	U
C21-TAS	U	U	U	U
C26-TAS(20S)	U	U	U	U
C26,C27-TAS	U	388.11 J	368.9 J	576.43 J
C27-TAS(20R)	U	226.69 J	U	454.53 J
C28-TAS(20S)	U	U	U	227.67 J
C28-TAS(20R)	Ŭ	Ŭ	Ŭ	261.58 J
C21-MAS	Ŭ	Ŭ	Ŭ	U
C22-MAS	Ŭ	Ŭ	Ŭ	U
C27-MAS	Ŭ	Ŭ	Ŭ	U
C27-20R-MAS	U	U	U	U
C27-20S-MAS	U	U	U	U
C28-20S-MAS	U	U	U	U
C27-C2920S/R-MAS	U	U	U	U
C29-20S-MAS	U	U	U	U
C29-20R-MAS	U	U	U	U
TAS_245	NA	NA	NA	NA
MAS_239	NA	NA	NA	NA

Surrogate Recoveries (%)				
5b(H)-Cholane	155388 NME	153337 NME	170266 NME	19515 NME

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07T09	GWP07T10	GWP07T11	GWP07T12
Battelle ID	R5532-P	R5533-P	R5534-P	R5535-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/11/07	01/11/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/08/07	02/09/07	02/11/07	02/08/07
Analytical Instrument	MS	MS	MS	MS
% Moisture	19.46	32.51	5.51	10.85
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.72	1.45	2.05	1.79
Size Unit-Basis	G DRY	G DRY	G DRY	G DRY
Minimum Reporting Limit	1080.96	699.41	979.51	415.48
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
C23 Tricyclic Terpane	U	U	DU	U
C29 Tricyclic Terpane -22S	U	U	DU	U
C29 Tricyclic Terpane -22R	U	U	DU	U
18a(H)-22,29,30-Trisnorneohopane -TS	U	U	DU	U
17a(H)-22,29,30-Trisnorhopane -TM	U	U	DU	U
30-Norhopane	U	U	DU	U
18a(H) & 18b(H)-Oleananes	U	U	DU	U
Hopane	U	U	DU	U
30-Homohopane -22S	U	U	DU	U
30-Homohopane -22R	U	U	DU	U
13b(H),17a(H)-20S-Diacholestane	U	U	DU	U
13b(H),17a(H)-20R-Diacholestane	U	U	DU	U
14a(H),17a(H)-20R-methylcholestane	U	U	3464.38 D	U
14a(H),17a(H)-20S-Ethylcholestane	U U	U	1275.21 DJ	U U
14a(H),17a(H)-20R-Ethylcholestane		U U	2158.71 D	UU
C21-TAS	U	-	DU	UU
C26-TAS(20S) C26,C27-TAS	U 646.21 J	159.25 J 392.28 J	510.92 DJ 1941.12 D	170.77 J
C27-TAS(20R)	469.54 J	233.02 J	1941.12 D 1115.3 D	100.81 J
C28-TAS(20S)	409.34 J 304.97 J	148.22 J	670.63 DJ	66.82 J
C28-TAS(208)	361.69 J	140.22 J U	690.6 DJ	81.1 J
C21-MAS	U	U	DU	U
C22-MAS	U	Ű	DU	U
C27-MAS	U	Ŭ	DU	Ű
C27-20R-MAS	Ŭ	Ŭ	DU	Ŭ
C27-20S-MAS	Ŭ	Ŭ	DU	Ŭ
C28-20S-MAS	Ŭ	Ŭ	DU	Ŭ
C27-C2920S/R-MAS	Ŭ	Ŭ	DU	Ŭ
C29-20S-MAS	U	U	DU	Ŭ
C29-20R-MAS	U	Ŭ	DU	Ŭ
TAS_245	NA	NA	NA	NA
MAS_239	NA	NA	NA	NA
Surrogate Recoveries (%)				
U				

5b(H)-Cholane	32311 NME	27454 NME	5507 ND	48227 NME

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07T13	GWP07T14	GWP07S01	GWP07S02
Battelle ID	R5536-P	R5537-P	R5538-P	R5539-P
Sample Type	SA	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/12/07	01/12/07
Extraction Date	01/24/07	01/24/07	01/24/07	01/24/07
Analysis Date	02/09/07	02/09/07	02/09/07	02/08/07
Analytical Instrument	MS	MS	MS	MS
% Moisture	12.18	7.01	13.85	10.23
% Lipid	NA	NA	NA	NA
Matrix	TAR	TAR	TAR	TAR
Sample Size	1.83	1.95	1.75	1.80
Size Unit-Basis	G_DRY	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit	385.01	12.94	1006.52	1032.92
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
Onito	NG/G_BIT		NG/G_DICI	NO/O_DICI
C23 Tricyclic Terpane	U	82.07	U	U
C29 Tricyclic Terpane -22S	Ŭ	55.09	Ŭ	Ŭ
C29 Tricyclic Terpane -22R	Ŭ	42.19	Ŭ	U
18a(H)-22,29,30-Trisnorneohopane -TS	Ŭ	33.26	Ŭ	Ŭ
17a(H)-22,29,30-Trisnorhopane -TM	Ŭ	50.17	Ŭ	Ŭ
30-Norhopane	Ŭ	167.1	Ŭ	U
18a(H) & 18b(H)-Oleananes	U	92.31	U	U
Hopane	U	308.82	Ű	U
30-Homohopane -22S	U	99.1	U	U
30-Homohopane -22R	U	73.11	Ű	U
13b(H),17a(H)-20S-Diacholestane	U	190.73	Ű	U
13b(H),17a(H)-20R-Diacholestane	U	127.58	Ű	U
14a(H),17a(H)-20R-methylcholestane	U	187.19	U	U
14a(H),17a(H)-20S-Ethylcholestane	U	86.87	U	U
14a(H),17a(H)-20R-Ethylcholestane	U	154.99	Ű	U
C21-TAS	U	U	Ű	U
C26-TAS(20S)	U	Ŭ	Ű	U
C26,C27-TAS	127.76 J	45.4	U	U
C27-TAS(20R)	78.73 J	30.21	U	U
C28-TAS(20S)	218.3 J	20.81	Ű	U
C28-TAS(20R)	162.12 J	16.11	Ű	U
C21-MAS	U	U	U	U
C22-MAS	U	Ŭ	Ŭ	U
C27-MAS	Ŭ	Ŭ	Ŭ	Ŭ
C27-20R-MAS	U	Ŭ	21465.67	17342.52
C27-20S-MAS	Ŭ	Ŭ	3524.46	2631.01
C28-20S-MAS	Ŭ	Ŭ	U	U
C27-C2920S/R-MAS	Ŭ	Ŭ	Ŭ	Ŭ
C29-20S-MAS	U	Ŭ	Ű	U
C29-20R-MAS	U	Ŭ	Ű	U
TAS 245	NA	NA	NA	NA
MAS_239	NA	NA	NA	NA
_				

Surrogate Recoveries (%)				
5b(H)-Cholane	46836 NME	212 NME	46848 NME	42830 NME

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07S03	GWP07S04	TDW3-4.5
Battelle ID	R5540-P	R5541-P	R5542-P
Sample Type	SA	SA	SA
Collection Date	01/12/07	01/12/07	09/26/06
Extraction Date	01/24/07	01/24/07	03/26/00
Analysis Date	01/24/07	02/08/07	02/09/07
5	02/12/07 MS	02/08/07 MS	02/09/07 MS
Analytical Instrument	_	_	-
% Moisture	20.14	12.16	14.86
% Lipid	NA	NA	NA
Matrix	TAR	TAR	SOIL
Sample Size	1.66	1.85	17.17
Size Unit-Basis	G_DRY	G_DRY	G_DRY
Minimum Reporting Limit	1120.03	1340.01	6.5
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY
C23 Tricyclic Terpane	U	U	17.62 T
C29 Tricyclic Terpane -22S	Ŭ	Ŭ	UT
C29 Tricyclic Terpane -22R	Ŭ	Ŭ	UT
18a(H)-22,29,30-Trisnorneohopane -TS	Ŭ	Ŭ	UT
17a(H)-22,29,30-Trisnorhopane -TM	Ŭ	Ŭ	26.61 T
30-Norhopane	Ŭ	Ŭ	69.06 T
18a(H) & 18b(H)-Oleananes	Ŭ	Ŭ	55.34 T
Hopane	Ŭ	Ŭ	113.88 T
30-Homohopane -22S	Ŭ	U	49.49 T
30-Homohopane -22R	Ŭ	Ŭ	38.99 T
13b(H),17a(H)-20S-Diacholestane	Ŭ	Ŭ	39.08 T
13b(H),17a(H)-20R-Diacholestane	Ŭ	Ŭ	22.14 T
14a(H),17a(H)-20R-methylcholestane	Ŭ	Ŭ	75.52 T
14a(H),17a(H)-20S-Ethylcholestane	Ŭ	Ŭ	43.36 T
14a(H),17a(H)-20R-Ethylcholestane	Ŭ	Ŭ	61.76 T
C21-TAS	Ŭ	Ŭ	UT
C26-TAS(20S)	Ŭ	Ŭ	UT
C26,C27-TAS	Ŭ	Ŭ	56.03 T
C27-TAS(20R)	Ŭ	Ŭ	35.08 T
C28-TAS(20S)	Ŭ	U	23.85 T
C28-TAS(20R)	Ŭ	Ŭ	20.94 T
C21-MAS	Ŭ	Ŭ	UT
C22-MAS	Ŭ	Ŭ	UT
C27-MAS	Ŭ	Ŭ	UT
C27-20R-MAS	11737.95	Ŭ	UT
C27-20S-MAS	4918.55	Ŭ	UT
C28-20S-MAS	U	Ŭ	UT
C27-C2920S/R-MAS	Ŭ	Ŭ	UT
C29-20S-MAS	Ŭ	U	UT
C29-20R-MAS	Ŭ	U	UT
TAS_245	NA	NA	NA
MAS_239	NA	NA	NA
—			

Surrogate Recoveries (%)

5b(H)-Cholane	48709 NME	20141 NME	691 NME
			••••

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID Procedural Blank Battelle ID BJ939PB-P Sample Type PB Collection Date 01/24/07 Analysis Date 02/10/07 Analysis Date 02/10/07 Analytical Instrument MS % Moisture 19.52 % Lipid NA Matrix SOIL, TAR Sample Size 2.44 Size Unit-Basis G_DRY Winimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane U C29 Tricyclic Terpane -22S U C23 Tricyclic Terpane -22R U Ba(H) & 18b(H)-Oleananes U 04-Nornohopane U 030-Homohopane -22S U 04-Homohopane -22S U 030-Homohopane -22S U 044a(H), 17a(H)-20R-methylcholestane U 13b(H), 17a(H)-20R-methylcholestane U 14a(H), 17a(H)-20R-Tehylcholestane U 224-TAS(20S) U U22-TAS(20R) U
Sample Type PB Collection Date 01/24/07 Extraction Date 01/24/07 Analytical Instrument MS % Moisture 19.52 % Lipid NA Matrix SOIL, TAR Sample Size 2.44 Size Unit-Basis G_DRY Vinimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane U C29 Tricyclic Terpane -22S U C29 Tricyclic Terpane -22R U U8a(H)-22,29,30-Trisnorneohopane -TS U V30-Norhopane U Waith & 18b(H)-Oleananes U 40pane U V30-Homohopane -22S U V30-Homohopane -22S U V30-Homohopane -22S U V4q(H), 17a(H)-20S-Diacholestane U V4a(H), 17a(H)-20S-Ethylcholestane U V4a(H), 17a(H)-20S-Ethylcholestane U V26-C27-TAS U V27-TAS(20R) U V22-TAS(20S) U
Sample Type PB Collection Date 01/24/07 Extraction Date 01/24/07 Analytical Instrument MS % Moisture 19.52 % Lipid NA Matrix SOIL, TAR Sample Size 2.44 Size Unit-Basis G_DRY Vinimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane U C29 Tricyclic Terpane -22S U C29 Tricyclic Terpane -22R U U8a(H)-22,29,30-Trisnorneohopane -TS U V30-Norhopane U Waith & 18b(H)-Oleananes U 40pane U V30-Homohopane -22R U V30-Homohopane -22S U V30-Homohopane -22R U V4a(H), 17a(H)-20S-Diacholestane U V4a(H), 17a(H)-20S-Ethylcholestane U V4a(H), 17a(H)-20S-Ethylcholestane U V26-C27-TAS U V27-TAS(20R) U V27-TAS(20R) U
Collection Date 01/24/07 Extraction Date 01/24/07 Analysis Date 02/10/07 Analytical Instrument MS & Moisture 19.52 & Lipid NA Matrix SOIL, TAR Sample Size 2.44 Size Unit-Basis G_DRY Winimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane 228 C23 Tricyclic Terpane -228 U C29 Tricyclic Terpane -228 U C29 Tricyclic Terpane -228 U U38(H)-22,29,30-Trisnorneohopane -TS U 30-Norhopane U 30-Norhopane U 30-Homohopane -228 U
Extraction Date 01/24/07 Analysis Date 02/10/07 Analysis Date 02/10/07 Analysis Date 02/10/07 Analysis Date 19.52 & Lipid NA Matrix SOIL, TAR Sample Size 2.44 Size Unit-Basis G_DRY Minimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane U C23 Tricyclic Terpane -22R U C29 Tricyclic Terpane -22R U Na(H)-22,29,30-Trisnorneohopane -TS U 17a(H)-22,29,30-Trisnorneohopane -TS U 030-Norhopane U 030-Norhopane U 13b(H), 17a(H)-20S-Diacholestane U 13b(H), 17a(H)-20S-Diacholestane U 13b(H), 17a(H)-20R-Diacholestane U 14a(H), 17a(H)-20R-Ethylcholestane U 14a(H), 17a(H)-20R-Ethylcholestane U 14a(H), 17a(H)-20R-Ethylcholestane U 122-TAS U 122-TAS(20R) U
Analysis Date 02/10/07 Analytical Instrument MS % Moisture 19.52 % Lipid NA Matrix SOIL, TAR Sample Size 2.44 Size Unit-Basis G_DRY Winimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane U C29 Tricyclic Terpane -22S U C29 Tricyclic Terpane -22R U 18a(H)-22,29,30-Trisnorneohopane -TS U 17a(H)-22,29,30-Trisnorneohopane -TS U 17a(H)-22,29,30-Trisnorneohopane -TS U 00-Norhopane U 30-Norhopane U 30-Homohopane -22S U 30-Homohopane -22S U 30-Homohopane -22S U 30-Homohopane -22R U 13b(H),17a(H)-20R-Diacholestane U 14a(H),17a(H)-20R-methylcholestane U 14a(H),17a(H)-20R-methylcholestane U 124-TAS U 127-TAS(20R) U 128-TAS(20R)
Analytical Instrument MS % Moisture 19.52 % Lipid NA Matrix SOIL, TAR Sample Size 2.44 Size Unit-Basis G_DRY Minimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane -22R U C29 Tricyclic Terpane -22R U C29 Tricyclic Terpane -22R U C29 Tricyclic Terpane -22R U 17a(H)-22,29,30-Trisnorneohopane -TS U 17a(H)-22,29,30-Trisnorneohopane -TS U 30-Norhopane U 8a(H) & 18b(H)-Oleananes U 40a-Homohopane -22S U 30-Homohopane -22R U 13b(H), 17a(H)-20S-Diacholestane U 13b(H), 17a(H)-20R-Diacholestane U 14a(H), 17a(H)-20R-Ethylcholestane U 14a(H), 17a(H)-20R-Ethylcholestane U 121-TAS U 122-TAS(20R) U 128-TAS(20R) U 128-TAS(20R) U 128-TAS(20R) U 127-MAS U <t< td=""></t<>
% Moisture 19.52 % Lipid NA Matrix SOIL, TAR Sample Size 2.44 Size Unit-Basis G_DRY Winimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane U C29 Tricyclic Terpane -22R U C29 Tricyclic Terpane -22R U 17a(H)-22,29,30-Trisnorhopane -TS U 00-Norhopane U 30-Norhopane U 30-Homohopane -22S U 30-Homohopane -22R U 30-Homohopane -22R U 30-Homohopane -22R U 30-Homohopane -22R U 13b(H),17a(H)-20S-Diacholestane U 13b(H),17a(H)-20R-methylcholestane U U21-TAS U C26-TAS(20S) U C28-TAS(20R) U C27-TAS U C28-TAS(20R) U C27-MAS U C27-MAS U C27-20S-MAS U
% Lipid NA Matrix SOIL, TAR Sample Size 2.44 Size Unit-Basis G_DRY Winimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane U C23 Tricyclic Terpane -22R U C29 Tricyclic Terpane -22R U Ba(H)-22,29,30-Trisnorneohopane -TS U Take(H)-22,29,30-Trisnorneohopane -TM U 30-Norhopane U 30-Norhopane U 30-Norhopane U 30-Homohopane -22R U 13b(H),17a(H)-20R-methylcholestane U 144(H),17a(H)-20R-methylcholestane U 124-TAS U C26-TAS(20S) U C26-TAS(20S) U C28-TAS(20R) U C28-TAS(20R) U C27-MAS U C27-OR-MAS
Matrix SOIL, TAR Sample Size 2.44 Size Unit-Basis G_DRY Winimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane U C29 Tricyclic Terpane -22R U C29 Tricyclic Terpane -22R U 18a(H)-22,29,30-Trisnorneohopane -TS U 17a(H)-22,29,30-Trisnorneohopane -TM U 30-Norhopane U 30-Homohopane U 30-Homohopane U 30-Homohopane -22R U 13b(H),17a(H)-20S-Diacholestane U 144(H),17a(H)-20R-methylcholestane U 144(H),17a(H)-20R-methylcholestane U 124-TAS U 124-TAS U 127-TAS(20R) U 127-TAS(20R) U 128-TAS(20R) U <tr< td=""></tr<>
Sample Size 2.44 Size Unit-Basis G_DRY Winimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane U C29 Tricyclic Terpane -22S U C29 Tricyclic Terpane -22R U 18a(H)-22,29,30-Trisnorneohopane -TS U 30-Norhopane U 18a(H)-22,29,30-Trisnorneohopane -TM U 30-Norhopane U 18a(H)-22,29,30-Trisnorneohopane -TS U 30-Norhopane U 18a(H)-22,29,30-Trisnorhopane -TM U 30-Norhopane U 18a(H)-22,29,30-Trisnorhopane -TS U 30-Norhopane U 18a(H)-22,29,30-Trisnorhopane -TM U 30-Norhopane U 18a(H)-22,29,30-Trisnorhopane -TM U 30-Homohopane -22R U 30-Homohopane -22R U 30-Homohopane -22R U 13b(H),17a(H)-20R-Ethylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U 124-TAS U C27-TAS(20R) U C28-TAS(20R)
Size Unit-Basis G_DRY Winimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane U C29 Tricyclic Terpane -22S U C29 Tricyclic Terpane -22R U 18a(H)-22,29,30-Trisnorneohopane -TS U 17a(H)-22,29,30-Trisnorneohopane -TM U 30-Norhopane U 18a(H) & 18b(H)-Oleananes U 40pane U 30-Homohopane -22S U 30-Homohopane -22S U 30-Homohopane -22R U 13b(H),17a(H)-20R-Diacholestane U 13b(H),17a(H)-20R-Diacholestane U 14a(H),17a(H)-20R-bitylcholestane U 14a(H),17a(H)-20R-bitylcholestane U 126-TAS U 127-TAS U 128-TAS(20R) U 128-TAS(20R) U 128-TAS(20R) U 127-TAS U 127-TAS(20R) U 128-TAS(20R) U 127-TAS(20R) U
Minimum Reporting Limit 8.23 Jnits NG/G_DRY C23 Tricyclic Terpane U C29 Tricyclic Terpane -22S U C29 Tricyclic Terpane -22R U 18a(H)-22,29,30-Trisnorneohopane -TS U 17a(H)-22,29,30-Trisnorhopane -TM U 30-Norhopane U 18a(H) & 18b(H)-Oleananes U -lopane U 30-Homohopane -22S U 30-Homohopane -22R U 13b(H),17a(H)-20S-Diacholestane U 13b(H),17a(H)-20R-Diacholestane U 14a(H),17a(H)-20R-Ethylcholestane U U22-TAS U C24-TAS U C26-TAS(20S) U C27-TAS U C28-TAS(20R) U C28-TAS(20R) U C22-MAS U C27-OR-MAS U C27-20S-MAS U C28-TAS(20S) U C27-20S-MAS U C27-20S-MAS U C28-20S-MAS U
Units NG/G_DRY C23 Tricyclic Terpane L C29 Tricyclic Terpane -22S L C29 Tricyclic Terpane -22R L C29 Tricyclic Terpane -22R L Ba(H)-22,29,30-Trisnorneohopane -TS L 17a(H)-22,29,30-Trisnorhopane -TM L 30-Norhopane L 8a(H) & 18b(H)-Oleananes L 40pane L 30-Homohopane -22S L 30-Homohopane -22R L 13b(H),17a(H)-20S-Diacholestane L 13b(H),17a(H)-20R-methylcholestane L 14a(H),17a(H)-20R-methylcholestane L 14a(H),17a(H)-20R-methylcholestane L 124-TAS L C26-TAS(20S) L C26-TAS(20S) L C27-TAS L C27-TAS L C28-TAS(20R) L C27-MAS L C27-OR-MAS L C27-OS-MAS L C27-20S-MAS L C28-20S/MAS L C28-20S/MAS
C23 Tricyclic Terpane L C29 Tricyclic Terpane -22S L C29 Tricyclic Terpane -22R L C29 Tricyclic Terpane -22R L Ba(H)-22,29,30-Trisnorneohopane -TS L 30-Norhopane L 8a(H)-22,29,30-Trisnorhopane -TM L 30-Norhopane L 8a(H) & 18b(H)-Oleananes L 4opane L 30-Homohopane -22S L 30-Homohopane -22R L 13b(H),17a(H)-20S-Diacholestane L 13b(H),17a(H)-20R-methylcholestane L 14a(H),17a(H)-20S-Ethylcholestane L 14a(H),17a(H)-20S-Ethylcholestane L 124-TAS L C26-TAS(20S) L C27-TAS L C28-TAS(20S) L C28-TAS(20R) L C27-MAS L C27-MAS L C27-OR-MAS L C27-20S-MAS L C28-COS/MAS L C27-2020S/R-MAS L C28-20S-MAS L C28-20S-MAS L <t< td=""></t<>
C29 Tricyclic Terpane -22S U C29 Tricyclic Terpane -22R U I8a(H)-22,29,30-Trisnorneohopane -TS U I7a(H)-22,29,30-Trisnorhopane -TM U 30-Norhopane U 88(H) & 18b(H)-Oleananes U 4opane U 30-Homohopane -22S U 30-Homohopane -22R U 214a(H),17a(H)-20R-methylcholestane U C28-TAS(20S) U C28-TAS(20R) U C28-TAS(20R)<
C29 Tricyclic Terpane -22S U C29 Tricyclic Terpane -22R U I8a(H)-22,29,30-Trisnorneohopane -TS U J7a(H)-22,29,30-Trisnorhopane -TM U 30-Norhopane U 30-Norhopane U 30-Norhopane U 30-Norhopane U 30-Homohopane -22S U 30-Homohopane -22R U 214a(H),17a(H)-20R-methylcholestane U C28-TAS(20S) U C28-TAS(20R)
C29 Tricyclic Terpane -22R U 18a(H)-22,29,30-Trisnorneohopane -TS U 17a(H)-22,29,30-Trisnorhopane -TM U 30-Norhopane U 18a(H) & 18b(H)-Oleananes U 4opane U 30-Homohopane -22S U 30-Homohopane -22R U 13b(H),17a(H)-20S-Diacholestane U 14a(H),17a(H)-20R-methylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U 124a(H),17a(H)-20R-Ethylcholestane U 124a(H),17a(H)-20R-Ethylcholestane U 124a(H),17a(H)-20R-Ethylcholestane U 122-TAS U 122-TAS U 122-TAS(20R) U 122-TAS(20R) U 122-TAS(20R) U 12
C29 Tricyclic Terpane -22R U 18a(H)-22,29,30-Trisnorneohopane -TS U 17a(H)-22,29,30-Trisnorhopane -TM U 30-Norhopane U 18a(H) & 18b(H)-Oleananes U 4opane U 30-Homohopane -22S U 30-Homohopane -22R U 13b(H),17a(H)-20S-Diacholestane U 14a(H),17a(H)-20R-methylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U 124-TAS U C27-TAS U C28-TAS(20S) U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C22-MAS U C22-MAS U C22-
18a(H)-22,29,30-Trisnorneohopane -TS U 17a(H)-22,29,30-Trisnorhopane -TM U 30-Norhopane U 18a(H) & 18b(H)-Oleananes U Hopane U 30-Homohopane -22S U 30-Homohopane -22R U 13b(H),17a(H)-20S-Diacholestane U 13b(H),17a(H)-20R-Diacholestane U 14a(H),17a(H)-20R-methylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U 121-TAS U C26-TAS(20S) U C26-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C27-MAS U C27-MAS U C27-MAS U C27-MAS U C27-20S-MAS U C28-CS-MAS U C28-20S/MAS U C28-20S/MAS U C28-20S/MAS U C28-20S/MAS U C28-
17a(H)-22,29,30-Trisnorhopane -TM U 30-Norhopane U 8a(H) & 18b(H)-Oleananes U 4opane U 30-Homohopane -22S U 30-Homohopane -22R U 13b(H),17a(H)-20S-Diacholestane U 13b(H),17a(H)-20R-Diacholestane U 14a(H),17a(H)-20R-methylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U 121-TAS U 224-TAS U 226-TAS(20S) U C26,C27-TAS U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C27-MAS U C27-MAS U C27-MAS U C27-20S-MAS U C28-CS-MAS U C28-CS-MAS U C28-20S-MAS U C28-20S-MAS U C28-20S-MAS U C28-20S-MAS U C28-20S-MAS U C29-20S-MAS U
30-Norhopane U 18a(H) & 18b(H)-Oleananes U 4opane U 30-Homohopane -22S U 30-Homohopane -22R U 13b(H), 17a(H)-20S-Diacholestane U 13b(H), 17a(H)-20R-Diacholestane U 13b(H), 17a(H)-20R-Diacholestane U 14a(H), 17a(H)-20R-Ethylcholestane U 14a(H), 17a(H)-20R-Ethylcholestane U 14a(H), 17a(H)-20R-Ethylcholestane U 126-TAS U 226-TAS(20S) U C26,C27-TAS U C28-TAS(20R) U C27-MAS U C27-WAS U C27-20S-MAS U C28-20S-MAS U C28-20S-MAS U C28-20S/R-MAS U C28-20S/MAS U
18a(H) & 18b(H)-Oleananes U Hopane U 30-Homohopane -22S U 30-Homohopane -22R U 13b(H),17a(H)-20S-Diacholestane U 13b(H),17a(H)-20S-Diacholestane U 13b(H),17a(H)-20R-methylcholestane U 14a(H),17a(H)-20R-methylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U 126-TAS(20S) U C26-TAS(20S) U C27-TAS(20R) U C28-TAS(20S) U C28-TAS(20S) U C24-TAS U C22-MAS U C22-MAS U C22-MAS U C22-MAS U C27-Z90S-MAS U C28-20S/R-MAS U C28-20S-MAS U C28-20
Hopane U 30-Homohopane -22S U 30-Homohopane -22R U 13b(H),17a(H)-20S-Diacholestane U 14a(H),17a(H)-20R-methylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U C21-TAS U C26-TAS(20S) U C26-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C21-MAS U C22-MAS U C22-MAS U C27-20R-MAS U C27-20S-MAS U C28-TAS(20S) U C27-C2920S/R-MAS U C27-202-MAS U C27-220S-MAS U C28-20S-MAS U C28-20S-MAS U
30-Homohopane -22S U 30-Homohopane -22R U 30-Homohopane -22R U 30-Homohopane -22R U 30-Homohopane -22R U 13b(H),17a(H)-20S-Diacholestane U 13b(H),17a(H)-20R-Diacholestane U 14a(H),17a(H)-20R-methylcholestane U 14a(H),17a(H)-20S-Ethylcholestane U 14a(H),17a(H)-20R-thylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U C21-TAS U C24-TAS(20S) U C26-TAS(20S) U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C21-MAS U C22-MAS U C27-20R-MAS U C27-20S-MAS U C28-TAS(20S/R-MAS U C28-20S-MAS U C28-20S-MAS U C28-20S-MAS U C28-20S-MAS U C28-20S-MAS U C29-20S-MAS U
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13b(H),17a(H)-20S-Diacholestane U 13b(H),17a(H)-20R-Diacholestane U 14a(H),17a(H)-20R-methylcholestane U 14a(H),17a(H)-20S-Ethylcholestane U 14a(H),17a(H)-20S-Ethylcholestane U 12b(H),17a(H)-20R-Ethylcholestane U 12ch-TAS U 221-TAS U 226-TAS(20S) U C26,C27-TAS U C28-TAS(20R) U C27-MAS U C27-MAS U C27-MAS U C27-20S-MAS U C28-20S-MAS U C28-20S/R-MAS U C28-20S/R-MAS U C29-20S/MAS U
13b(H),17a(H)-20R-Diacholestane U 14a(H),17a(H)-20R-methylcholestane U 14a(H),17a(H)-20S-Ethylcholestane U 14a(H),17a(H)-20S-Ethylcholestane U 124a(H),17a(H)-20R-Ethylcholestane U 124a(H),17a(H)-20R-Ethylcholestane U 021-TAS U 026-TAS(20S) U 026,C27-TAS U 027-TAS(20R) U 028-TAS(20S) U 028-TAS(20R) U 028-TAS(20R) U 022-MAS U 022-MAS U 027-Q2-MAS U 027-20S-MAS U 028-20S-MAS U 028-20S-MAS U 029-20S-MAS U 029-20S-MAS U 029-20S-MAS U 029-20S-MAS U
14a(H),17a(H)-20R-methylcholestane U 14a(H),17a(H)-20S-Ethylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U C21-TAS U C26-TAS(20S) U C26,C27-TAS U C27-TAS(20R) U C28-TAS(20S) U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C21-MAS U C22-MAS U C27-Z0R-MAS U C27-C20S-MAS U C27-C2920S/R-MAS U C27-C2920S/R-MAS U C29-20S-MAS U
14a(H),17a(H)-20S-Ethylcholestane U 14a(H),17a(H)-20R-Ethylcholestane U C21-TAS U C26-TAS(20S) U C26-C27-TAS U C27-TAS(20R) U C28-TAS(20S) U C28-TAS(20R) U C28-TAS(20R) U C22-MAS U C22-MAS U C27-QR-MAS U C27-20S-MAS U C28-20S-MAS U C28-20S-MAS U C29-20S-MAS U
14a(H),17a(H)-20R-Ethylcholestane U C21-TAS U C26-TAS(20S) U C26-TAS(20R) U C27-TAS(20R) U C28-TAS(20S) U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C21-MAS U C22-MAS U C27-20R-MAS U C27-20S-MAS U C28-C2920S/R-MAS U C29-20S-MAS U C29-20S-MAS U
C21-TAS U C26-TAS(20S) U C26,C27-TAS U C27-TAS(20R) U C28-TAS(20S) U C28-TAS(20R) U C27-MAS U C27-MAS U C27-20S-MAS U C28-20S-MAS U C28-20S/R-MAS U C29-20S-MAS U C29-20S-MAS U
C26-TAS(20S) U C26,C27-TAS U C27-TAS(20R) U C28-TAS(20S) U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C21-MAS U C22-MAS U C27-20R-MAS U C28-20S-MAS U C28-20S/R-MAS U C29-20S/R-MAS U C29-20S-MAS U
C26,C27-TAS U C27-TAS(20R) U C28-TAS(20S) U C28-TAS(20R) U C28-TAS(20R) U C28-TAS(20R) U C21-MAS U C22-MAS U C27-20R-MAS U C27-20S-MAS U C28-20S-MAS U C27-2020S/R-MAS U C29-20S-MAS U C29-20S-MAS U
C27-TAS(20R) U C28-TAS(20S) U C28-TAS(20R) U C21-MAS U C22-MAS U C27-MAS U C27-20R-MAS U C27-20S-MAS U C27-C2920S/R-MAS U C29-20S-MAS U C29-20S-MAS U C29-20S-MAS U
C28-TAS(20S) U C28-TAS(20R) U C21-MAS U C22-MAS U C27-MAS U C27-20R-MAS U C27-20S-MAS U C28-C29-20S-MAS U C29-20S-MAS U C29-20S-MAS U C29-20S-MAS U C29-20S-MAS U
C28-TAS(20R) U C21-MAS U C22-MAS U C27-MAS U C27-20R-MAS U C27-20S-MAS U C28-20S-MAS U C28-20S-MAS U C29-20S-MAS U C29-20S-MAS U C29-20S-MAS U
C21-MAS U C22-MAS U C27-MAS U C27-20R-MAS U C27-20S-MAS U C28-20S-MAS U C28-20S-MAS U C28-20S-MAS U C29-20S-MAS U C29-20S-MAS U
C22-MAS U C27-MAS U C27-20R-MAS U C27-20S-MAS U C28-20S-MAS U C27-C2920S/R-MAS U C29-20S-MAS U
C27-MAS U C27-20R-MAS U C27-20S-MAS U C28-20S-MAS U C27-C2920S/R-MAS U C29-20S-MAS U
C27-MAS U C27-20R-MAS U C27-20S-MAS U C28-20S-MAS U C27-C2920S/R-MAS U C29-20S-MAS U
C27-20R-MAS U C27-20S-MAS U C28-20S-MAS U C27-C2920S/R-MAS U C29-20S-MAS U
C27-20S-MAS U C28-20S-MAS U C27-C2920S/R-MAS U C29-20S-MAS U
C28-20S-MAS U C27-C2920S/R-MAS U C29-20S-MAS U
C27-C2920S/R-MAS U C29-20S-MAS U
C29-20S-MAS U
C29-20R-MAS U
TAS_245 NA
MAS_239 NA

Surrogate Recoveries (%)

5b(H)-Cholane

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

	060208-03: Sand,	
Client ID	White Quartz, -50+70	
Battelle ID	BJ940LCS-P	
Sample Type	LCS	
Collection Date	01/24/07	
Extraction Date	01/24/07	
Analysis Date	02/10/07	
Analytical Instrument	MS	
% Moisture	NA	
% Lipid	NA	
Matrix	SOIL, TAR	
Sample Size	20.01	
Size Unit-Basis	G_DRY	
Minimum Reporting Limit	1	
Units	NG/G_DRY	Target % Recovery Qualifier
C23 Tricyclic Terpane	U	
C29 Tricyclic Terpane -22S	Ŭ	
C29 Tricyclic Terpane -22R	U	
18a(H)-22,29,30-Trisnorneohopane -TS	Ŭ	
17a(H)-22,29,30-Trisnorhopane -TM	Ŭ	
30-Norhopane	Ŭ	
18a(H) & 18b(H)-Oleananes	Ŭ	
Hopane	U	
30-Homohopane -22S	Ŭ	
30-Homohopane -22R	Ū	
13b(H),17a(H)-20S-Diacholestane	Ŭ	
13b(H),17a(H)-20R-Diacholestane	U	
14a(H),17a(H)-20R-methylcholestane	U	
14a(H),17a(H)-20S-Ethylcholestane	U	
14a(H),17a(H)-20R-Ethylcholestane	U	
C21-TAS	U	
C26-TAS(20S)	U	
C26,C27-TAS	U	
C27-TAS(20R)	U	
C28-TAS(20S)	U	
C28-TAS(20R)	U	
C21-MAS	U	
C22-MAS	U	
C27-MAS	U	
C27-20R-MAS	U	
C27-20S-MAS	U	
C28-20S-MAS	U	
C27-C2920S/R-MAS	U	
C29-20S-MAS	U	
C29-20R-MAS	U	
TAS_245	NA	
MAS_239	NA	

Surrogate Recoveries (%)

5b(H)-Cholane

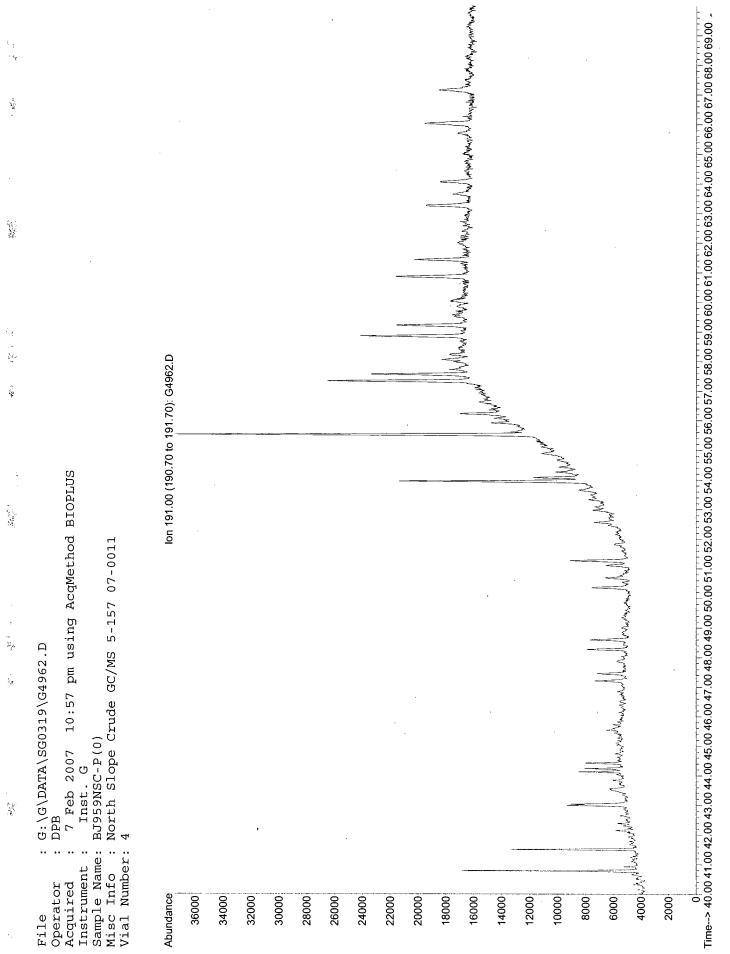
The Business of Innovation

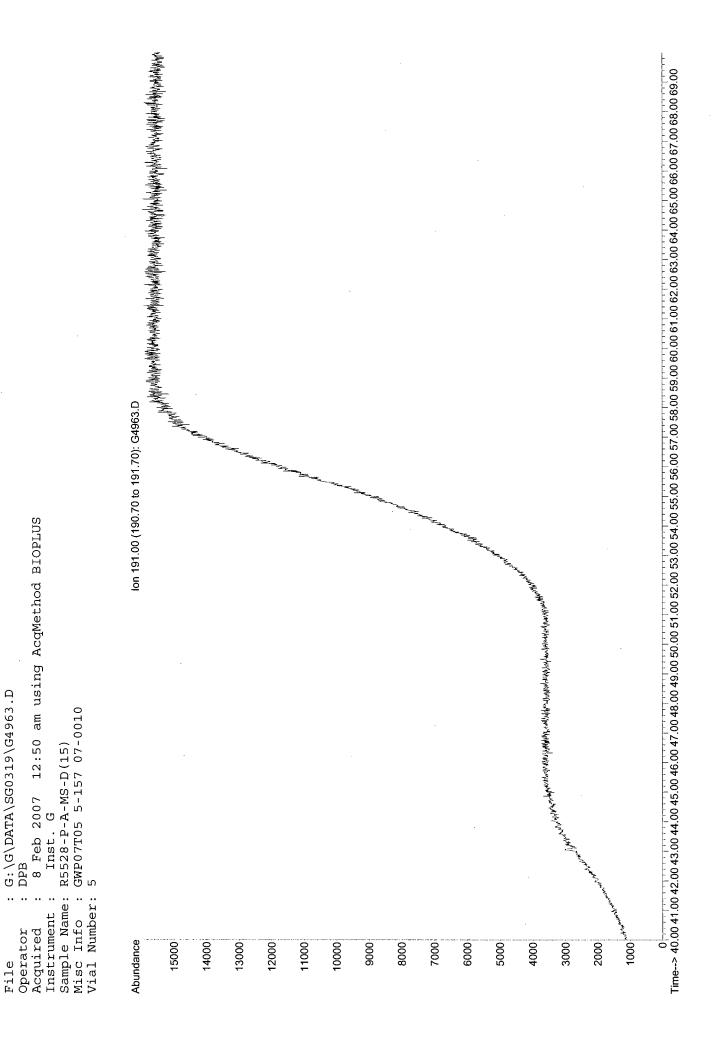
Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

	GN62: North Slope			
Client ID	Crude			
Cheft ID	Cidde			
Battelle ID	BJ959NSC-P			
Sample Type	NSC			
Collection Date	01/30/07			
Extraction Date	01/30/07			
Analysis Date	02/07/07			
Analytical Instrument	MS			
% Moisture	NA			
% Lipid	NA			
Matrix	OIL			
Sample Size	5.01			
Size Unit-Basis	G_OIL			
Minimum Reporting Limit	1.3			
Units	NG/G_OIL	Target % D	ifference	Qualifier
C23 Tricyclic Terpane	43.65	47.76	8.6	
C29 Tricyclic Terpane -22S	12.63	14.70	14.1	
C29 Tricyclic Terpane -22R	13.24	14.64	9.6	
18a(H)-22,29,30-Trisnorneohopane -TS	15.95	15.96	0.1	
17a(H)-22,29,30-Trisnorhopane -TM	24.18	24.82	2.6	
30-Norhopane	73.99	69.58	6.3	
18a(H) & 18b(H)-Oleananes	U	00100	0.0	
Hopane	124.37	120.14	3.5	
30-Homohopane -22S	62.52	59.93	4.3	
30-Homohopane -22R	38.83	39.69	2.2	
13b(H),17a(H)-20S-Diacholestane	44.55	44.18	0.8	
	25.35	25.52	0.8	
13b(H),17a(H)-20R-Diacholestane				
14a(H),17a(H)-20R-methylcholestane	35.49	33.94	4.6	
14a(H),17a(H)-20S-Ethylcholestane	41.61	35.93	15.8	
14a(H),17a(H)-20R-Ethylcholestane	40.28	39.17	2.8	
C21-TAS	18.59			
C26-TAS(20S)	15.53			
C26,C27-TAS	55.26			
C27-TAS(20R)	37.91			
C28-TAS(20S)	31.48			
C28-TAS(20R)	31.26			
C21-MAS	6.09			
C22-MAS	3.54			
C27-MAS	4.73			
C27-20R-MAS	5.98			
C27-20S-MAS	2.57			
C28-20S-MAS	14.9			
C27-C2920S/R-MAS	12.33			
C29-20S-MAS	4.08			
C29-20R-MAS	9.14			
TAS_245	NA			
MAS_239	NA			

Surrogate Recoveries (%)

5b(H)-Cholane

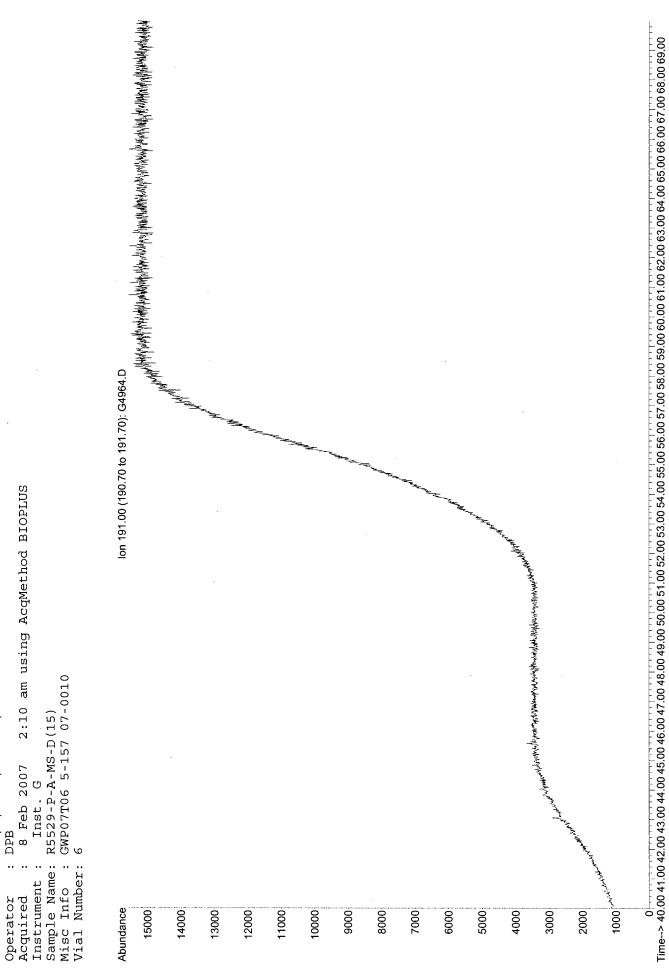




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A)

File

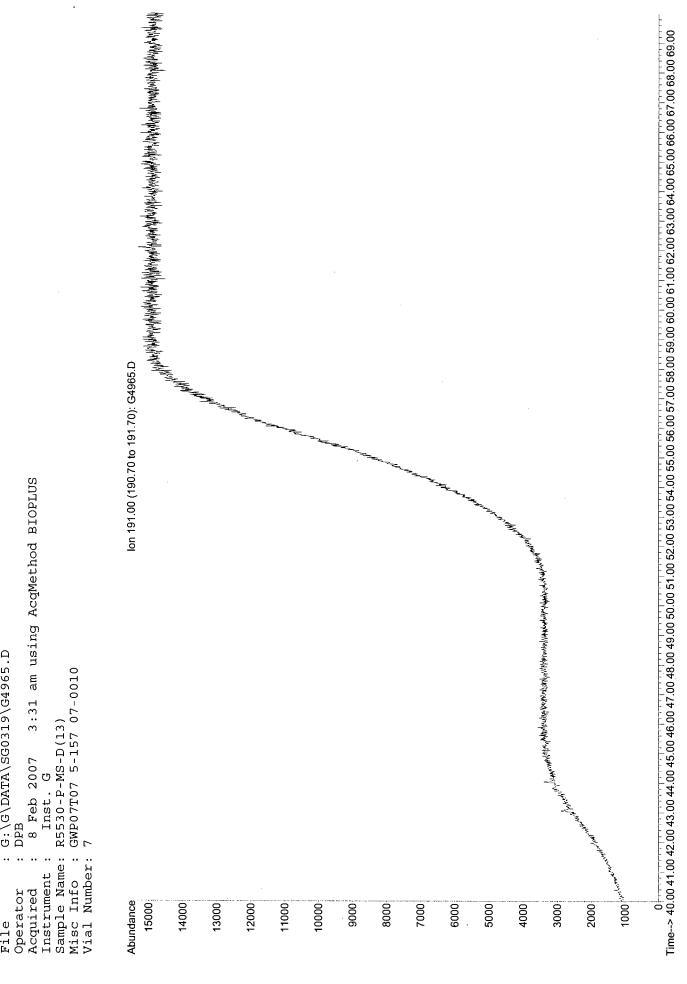


: G:\G\DATA\SG0319\G4964.D : DPB

File

4

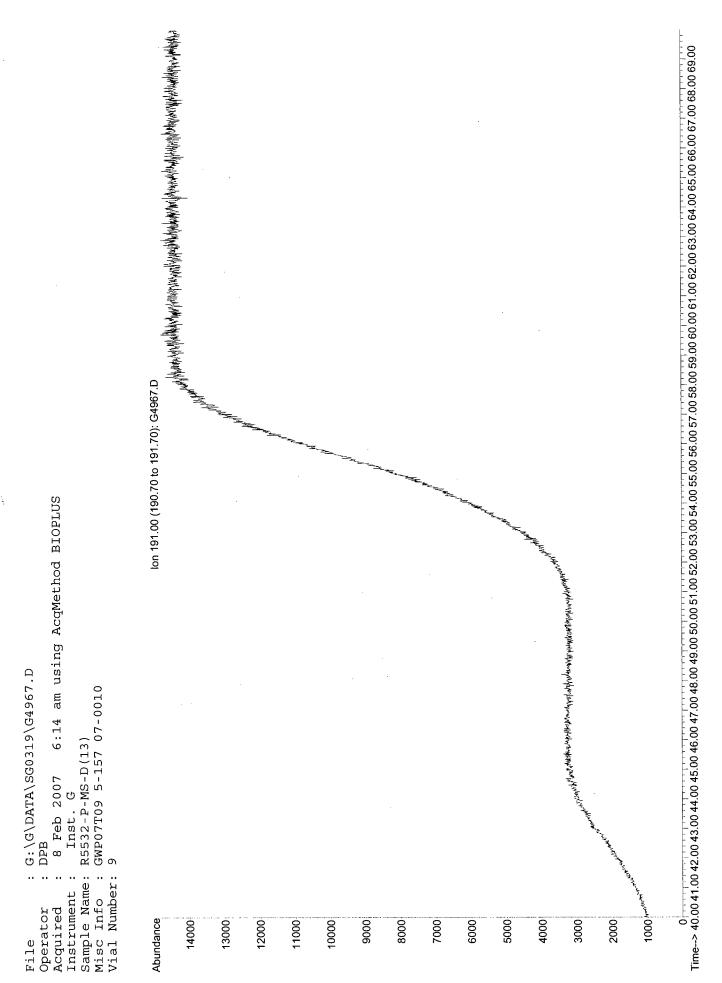
2:10 am using AcqMethod BIOPLUS



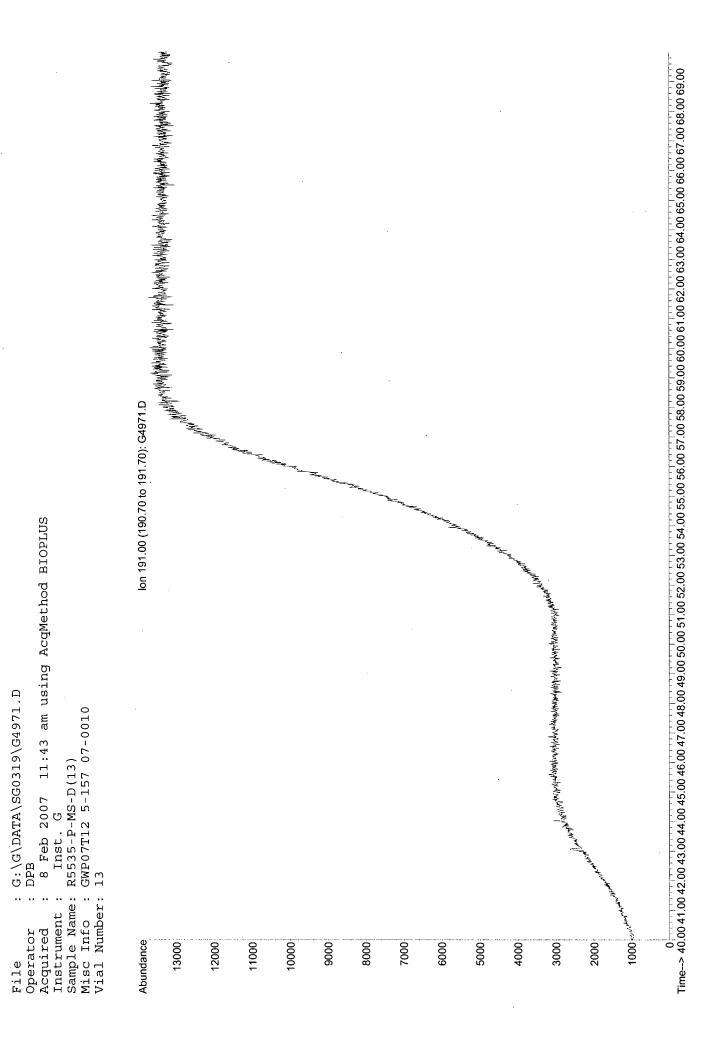
3:31 am using AcqMethod BIOPLUS

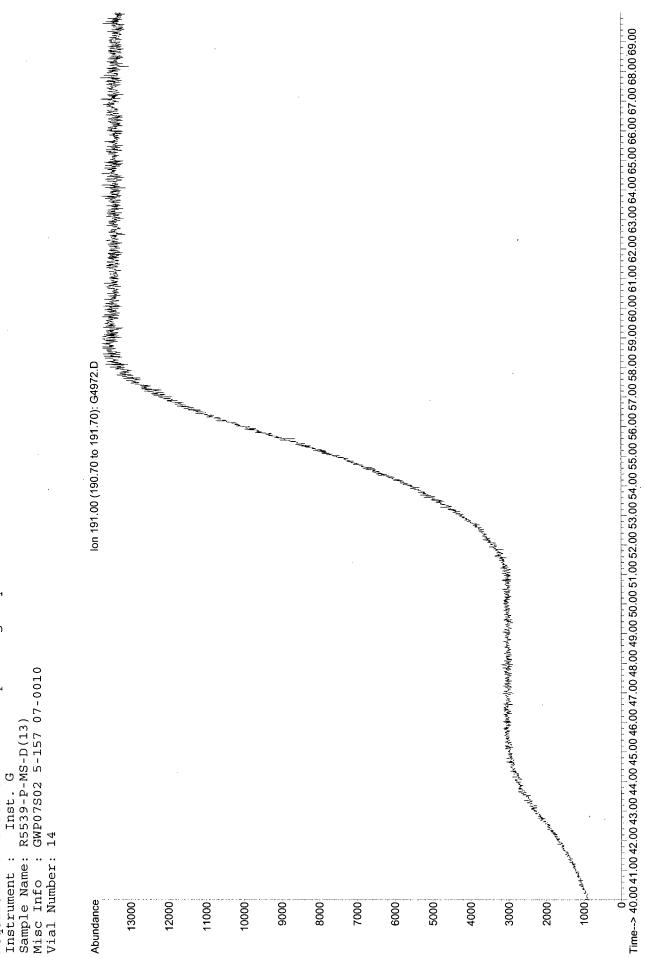
: G:\G\DATA\SG0319\G4965.D : DPB

File



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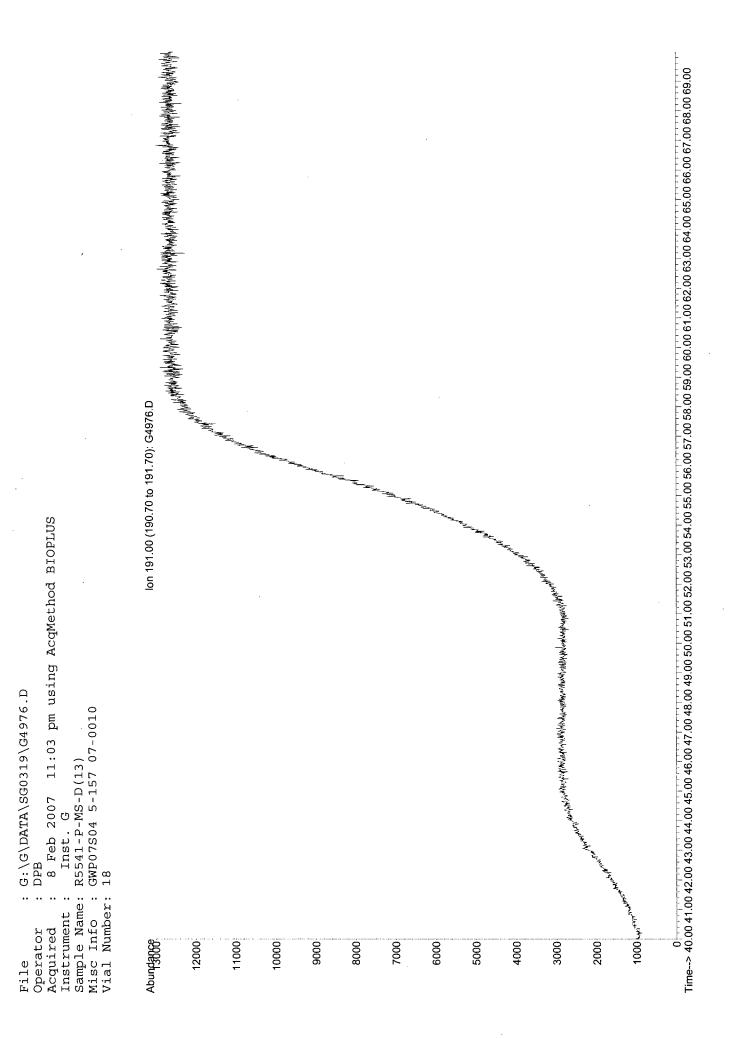


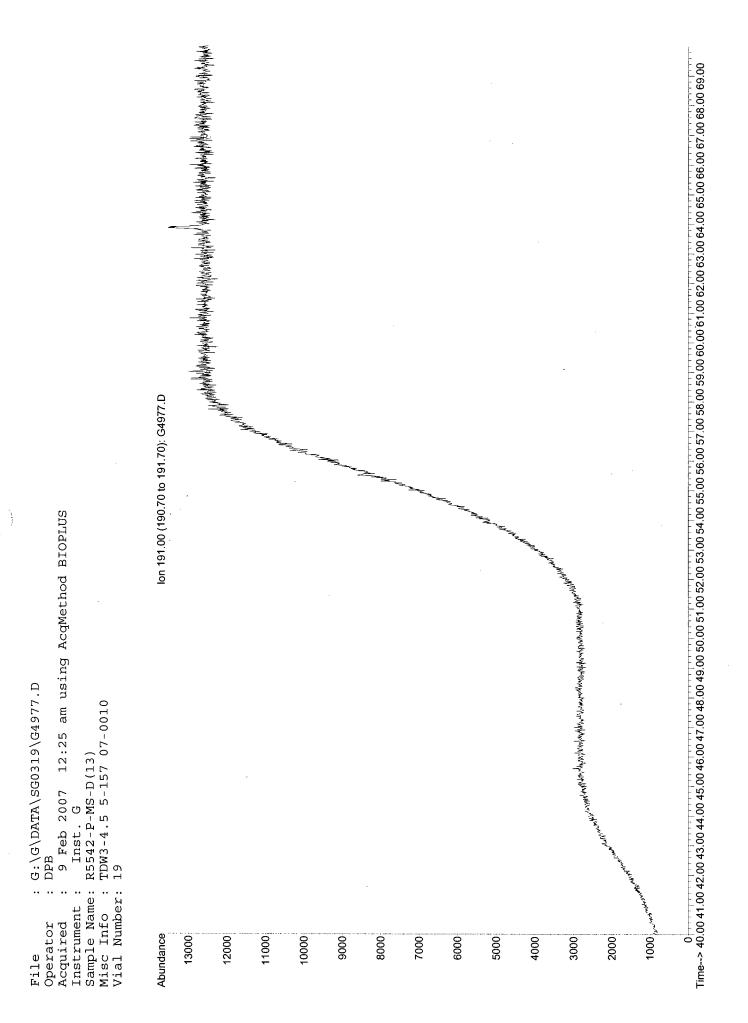


5:37 pm using AcqMethod BIOPLUS : G:\G\DATA\SG0319\G4972.D : DPB Acquired : 8 Feb 2007 5: Instrument : Inst. G Sample Name: R5539-P-MS-D(13) Operator

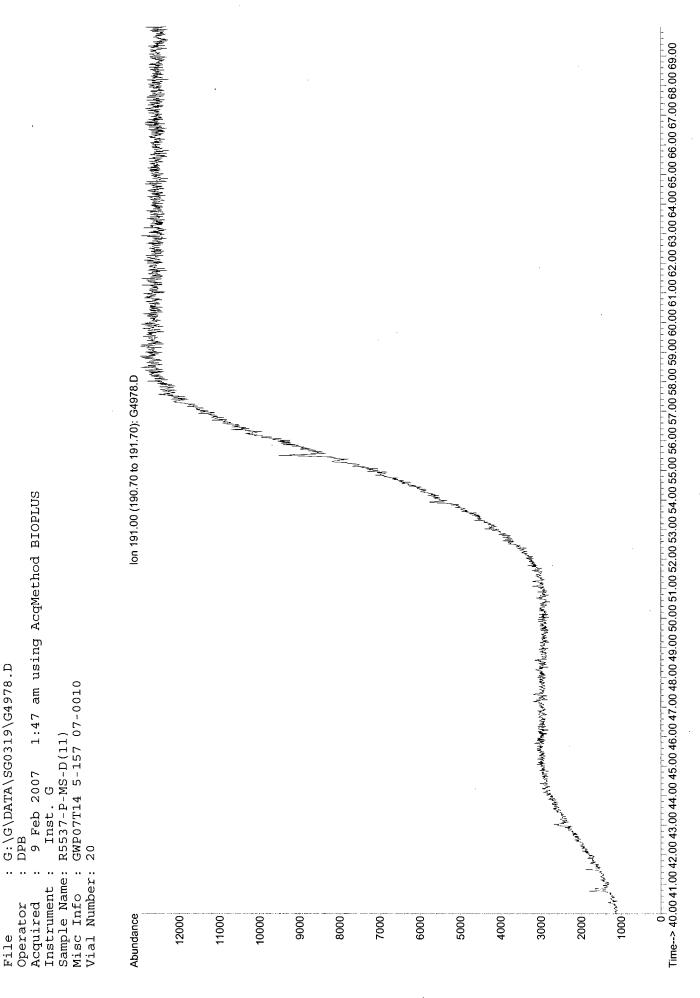
File

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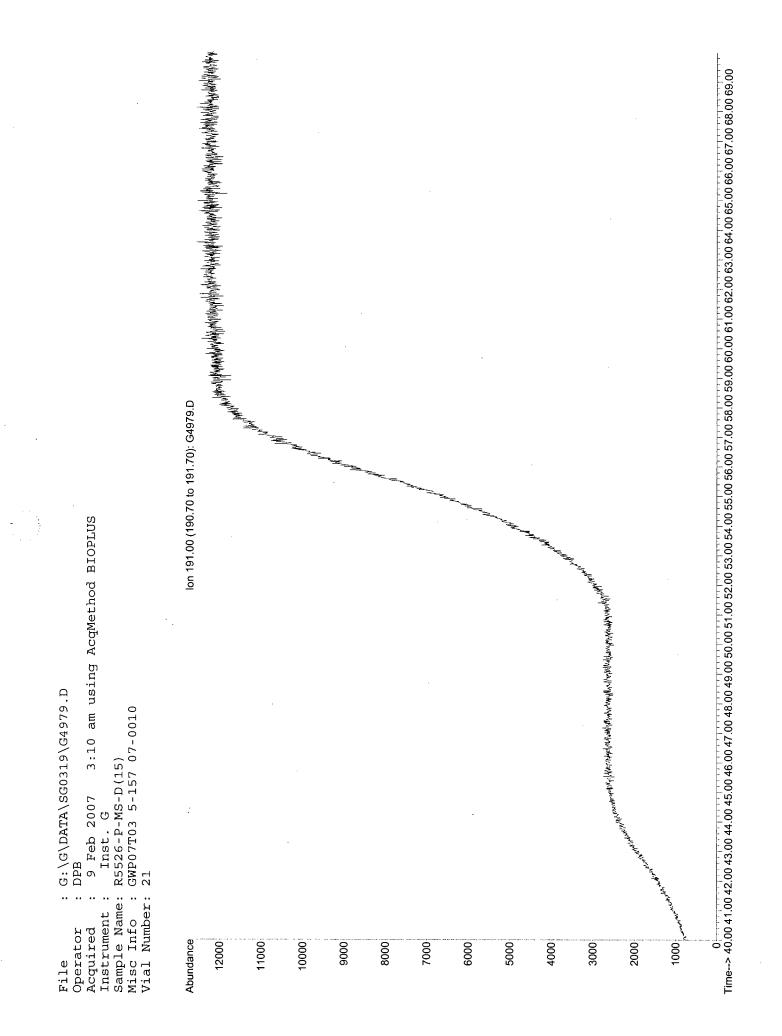


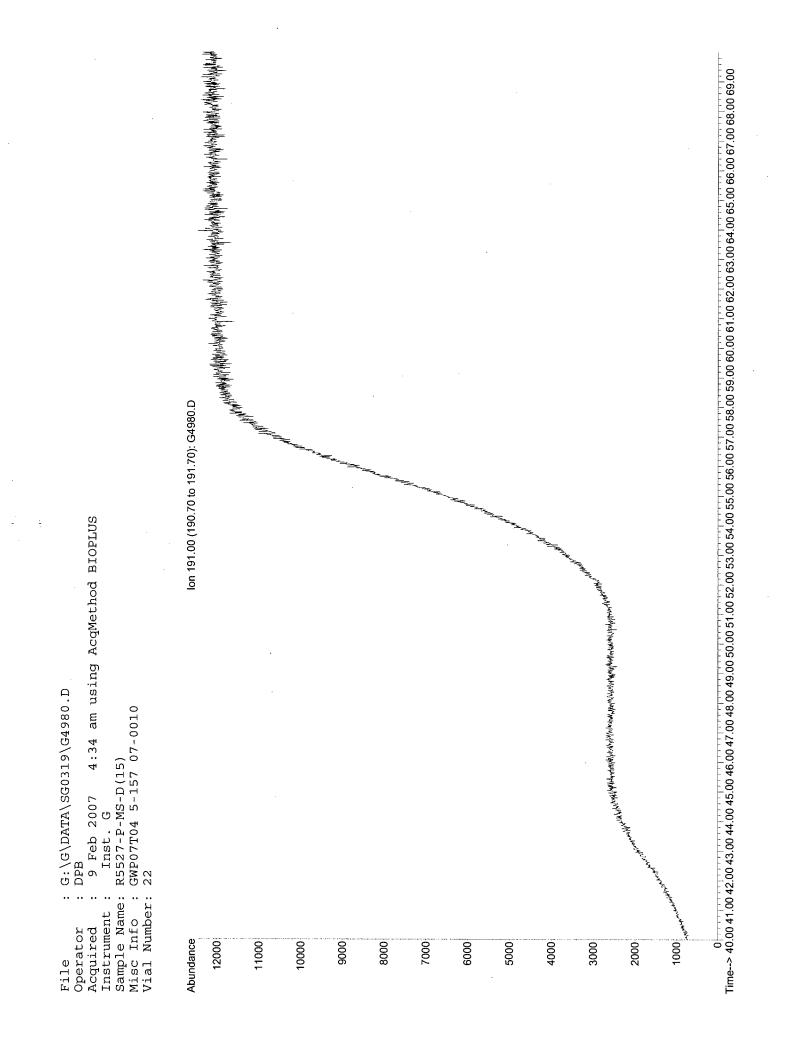


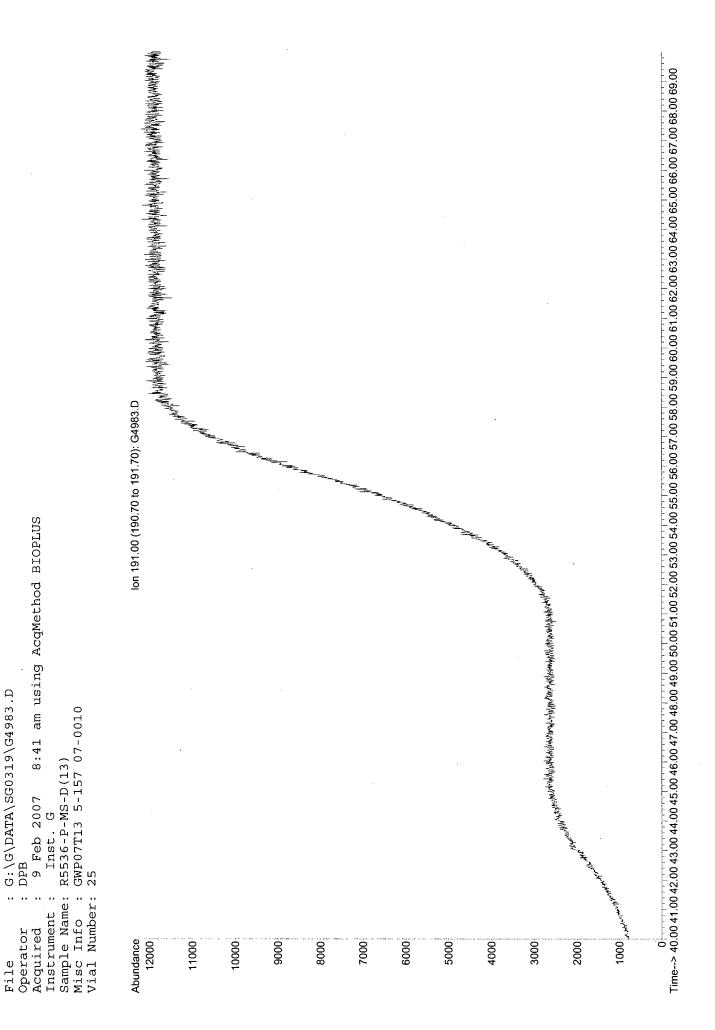
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1:47 am using AcqMethod BIOPLUS : G:\G\DATA\SG0319\G4978.D : DPB

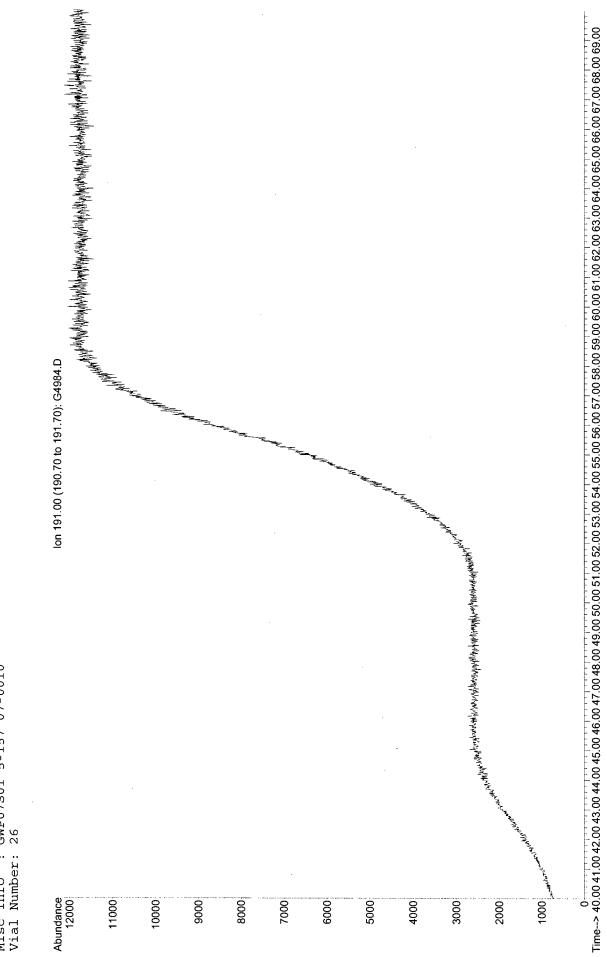




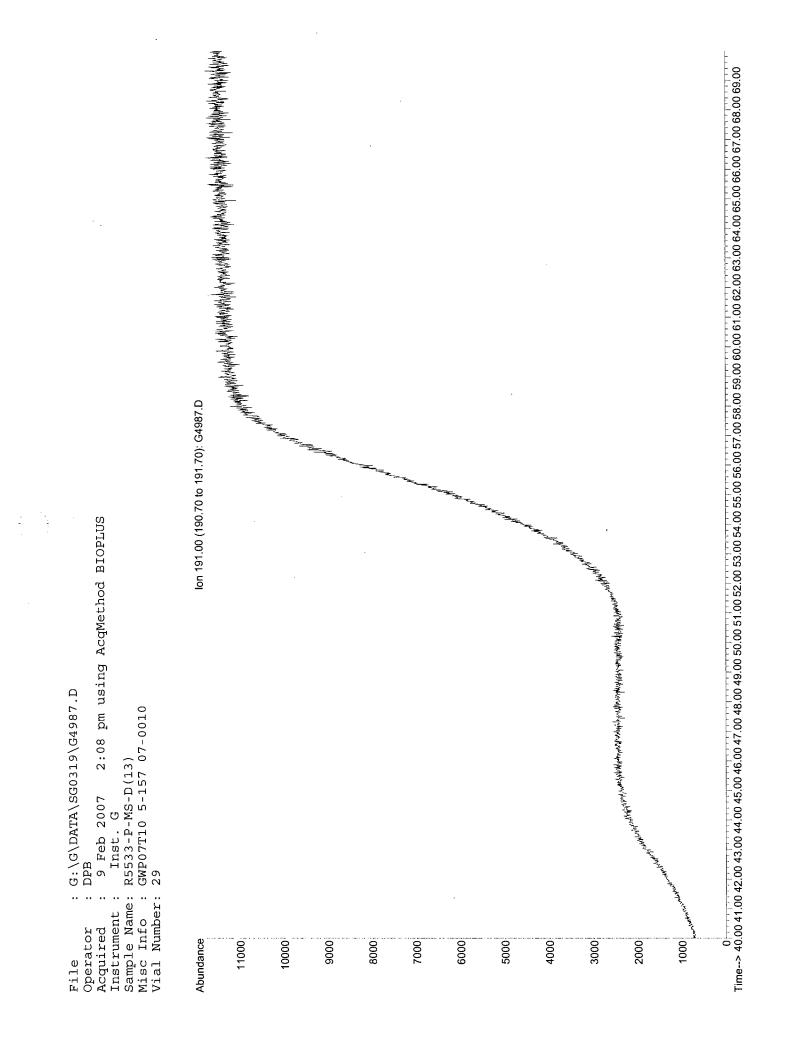


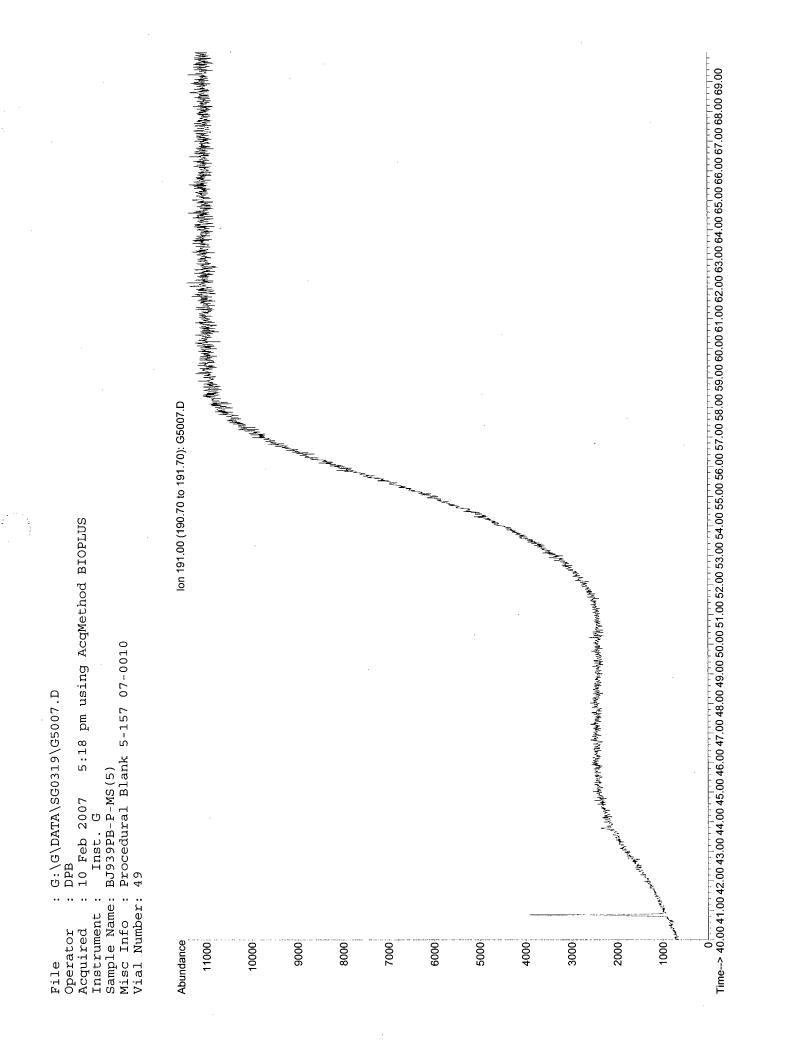
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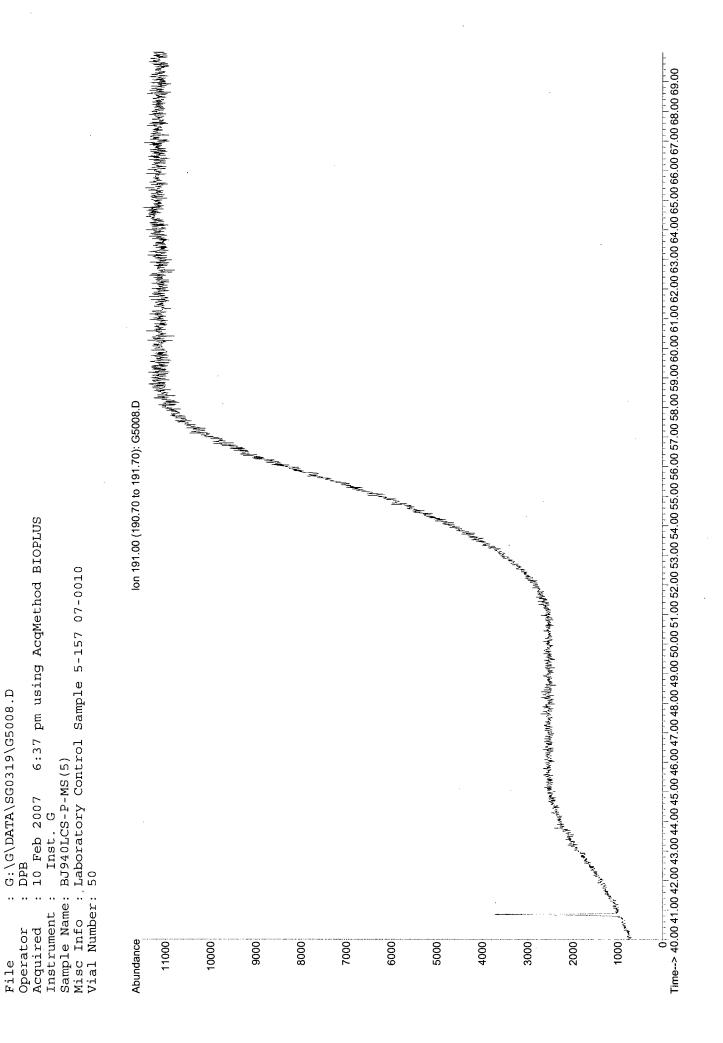
File

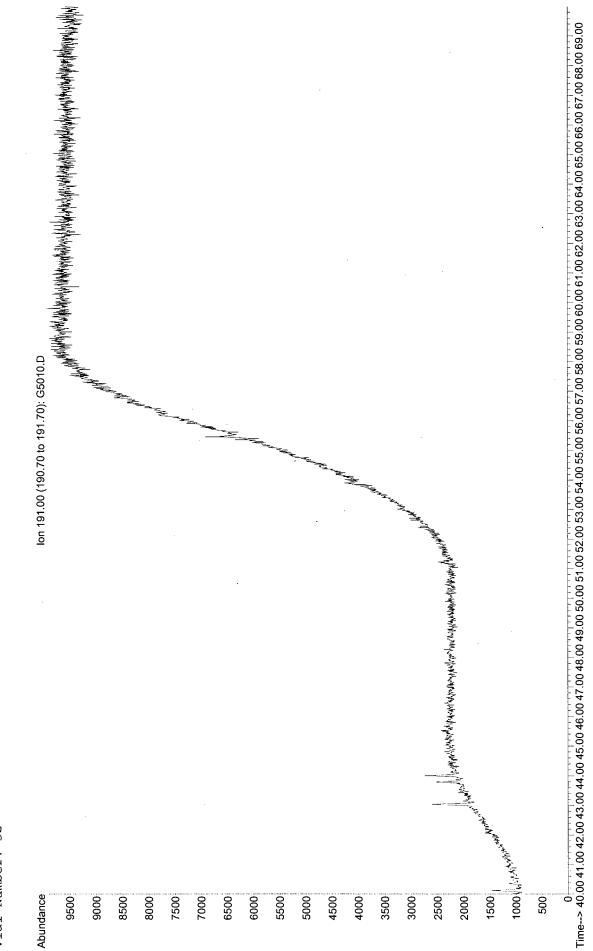


File : G:\G\DATA\SG0319\G4984.D Operator : DPB Acquired : 9 Feb 2007 10:03 am using AcqMethod BIOPLUS Instrument : Inst. G Sample Name: R5538-P-MS-D(13) Misc Info : GWP07S01 5-157 07-0010 Vial Number: 26

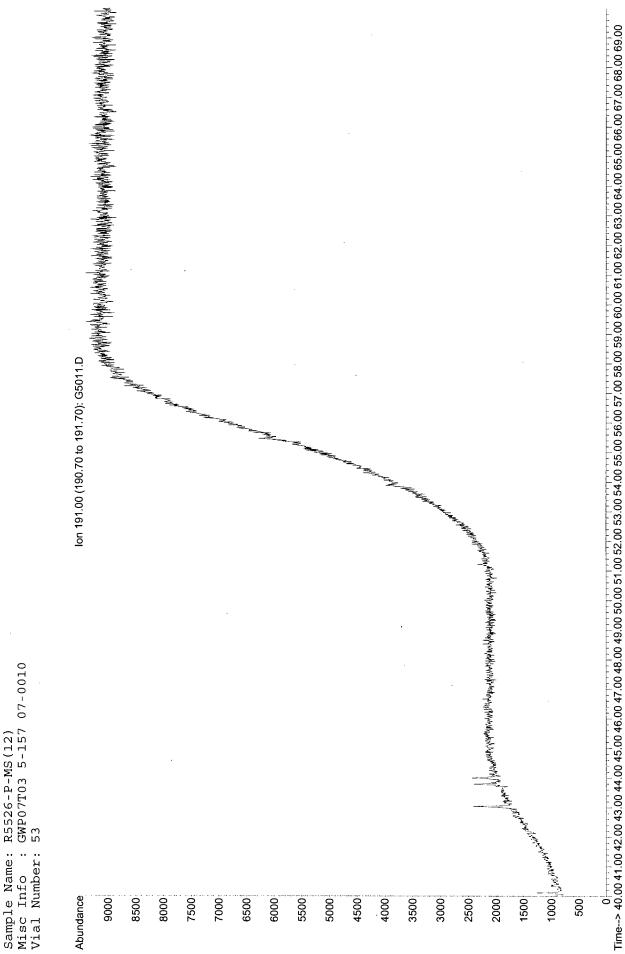






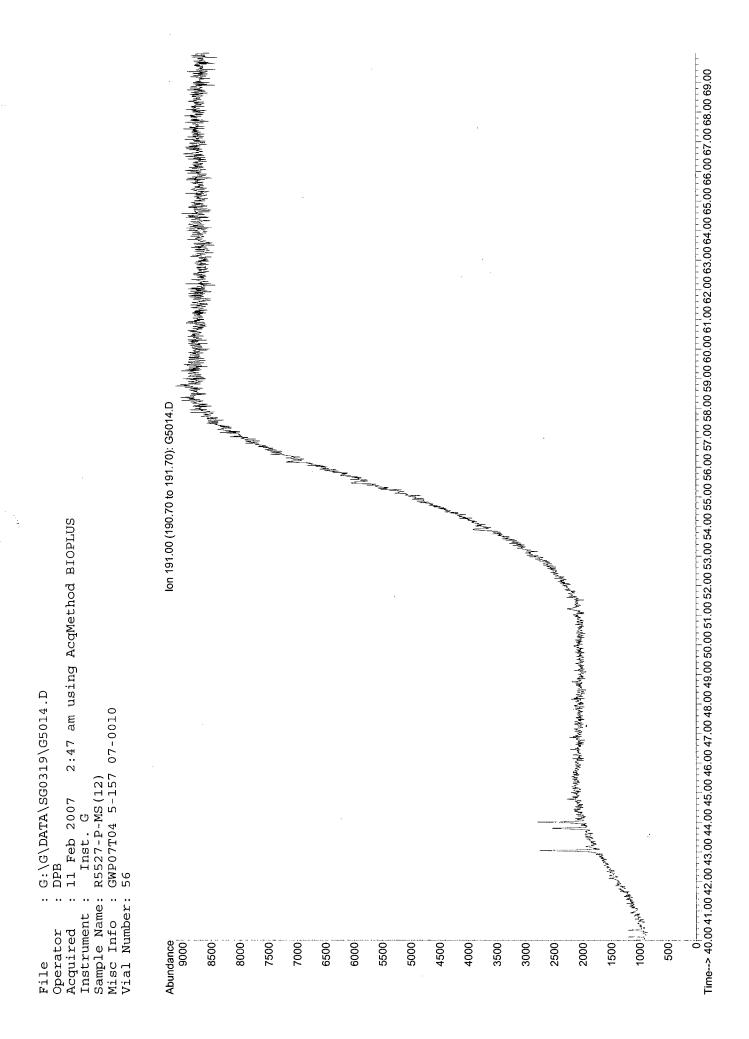


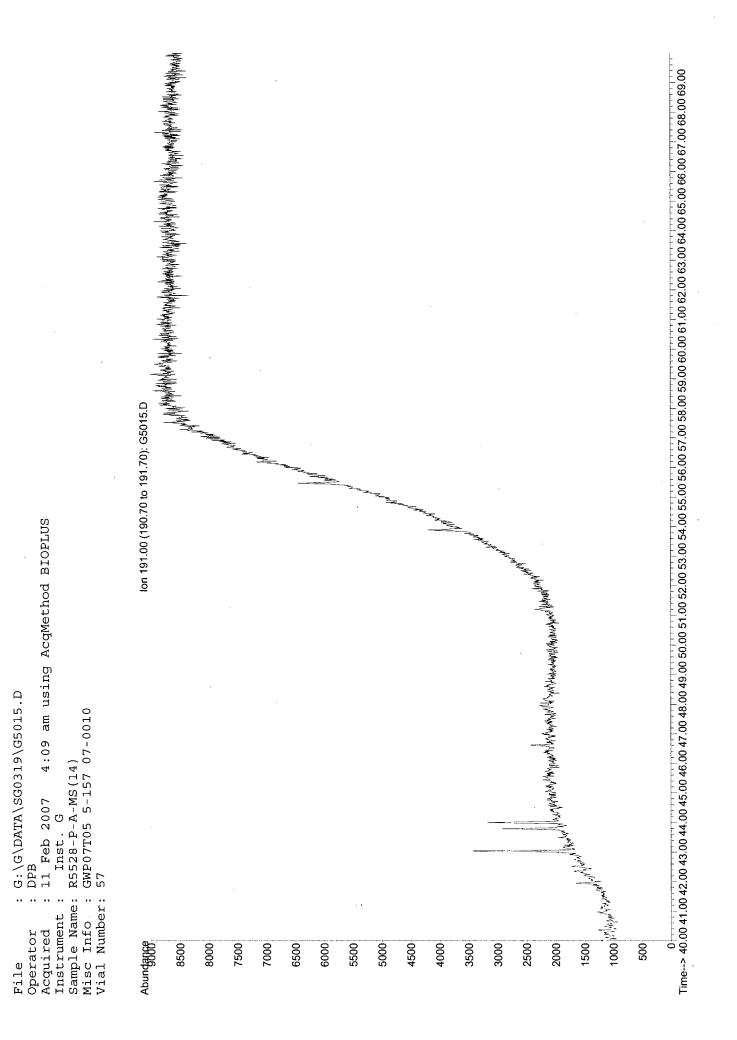
File : G:\G\DATA\SG0319\G5010.D Operator : DPB Acquired : 10 Feb 2007 9:21 pm using AcqMethod BIOPLUS Instrument : Inst. G Sample Name: R5525-P-MS(12) Misc Info : GWP07T02 5-157 07-0010 Vial Number: 52

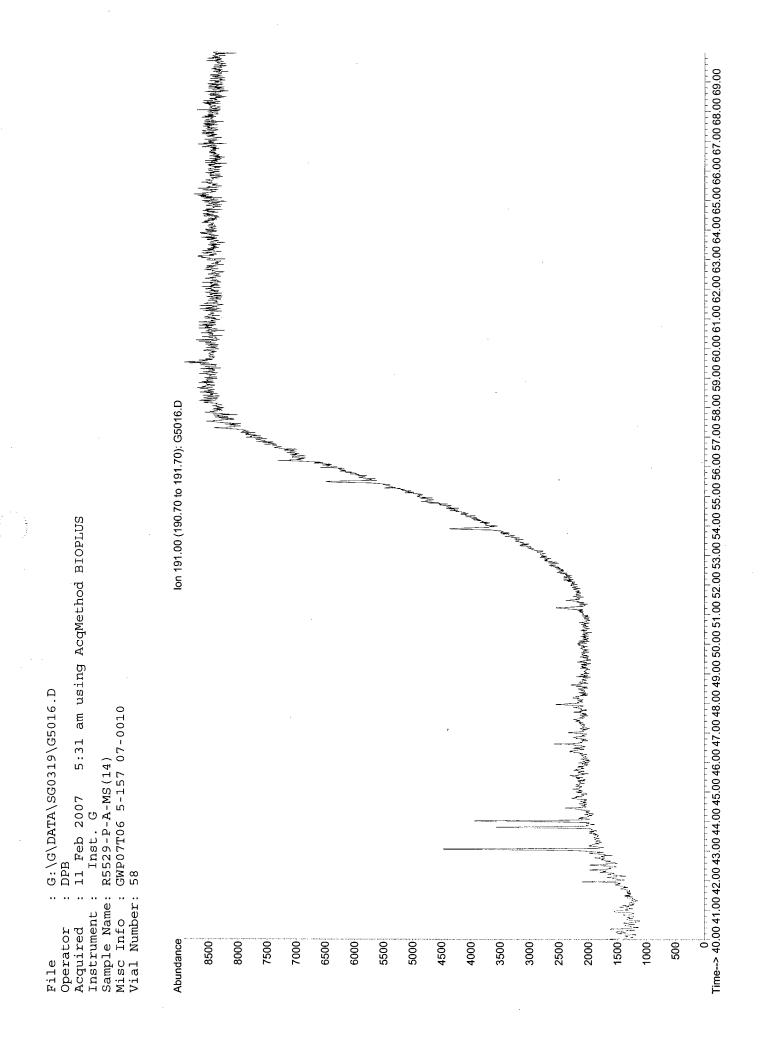


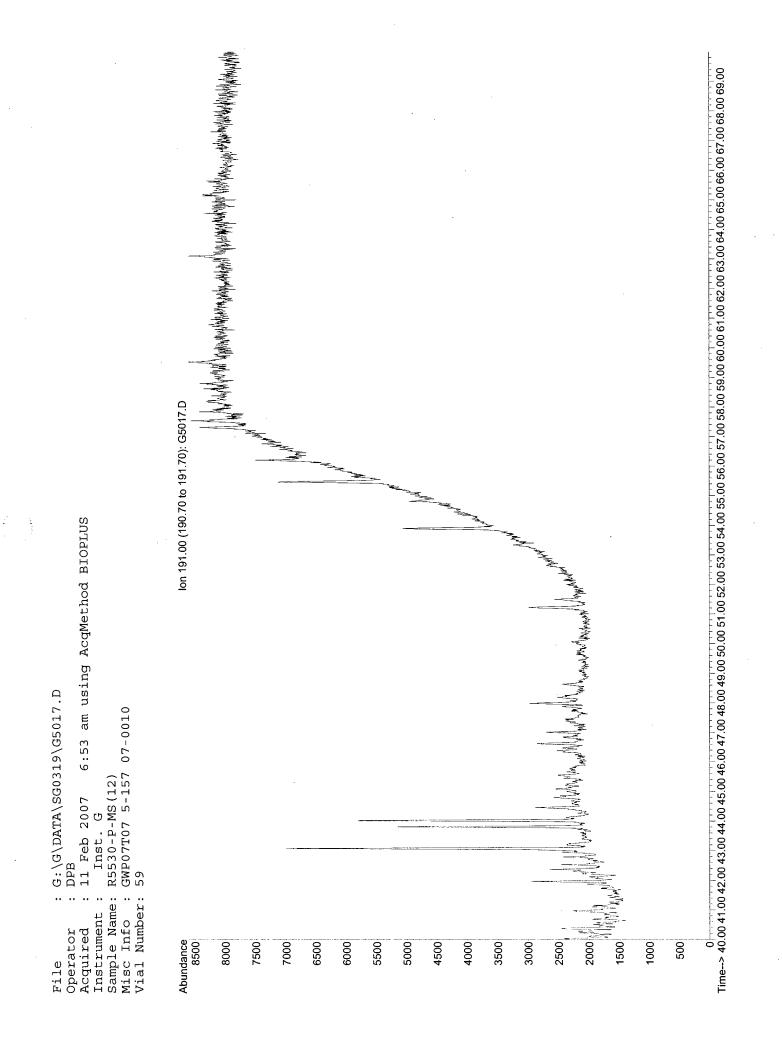
Acquired : 10 Feb 2007 10:43 pm using AcqMethod BIOPLUS Instrument : Inst. G Sample Name: R5526-P-MS(12) Misc Info : GWP07T03 5-157 07-0010 G:\G\DATA\SG0319\G5011.D DPB Operator File

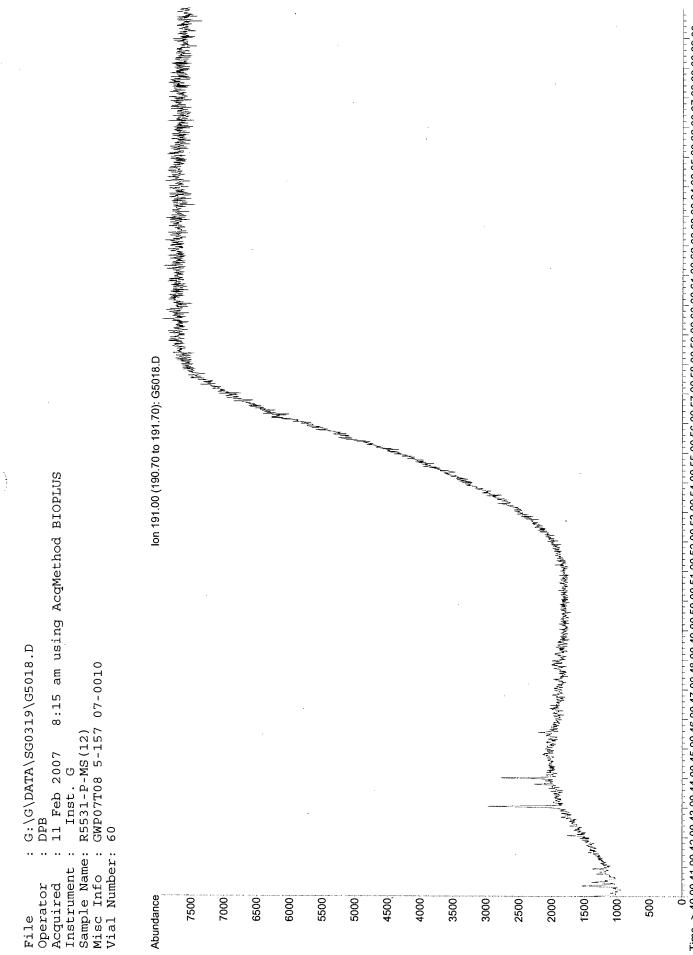
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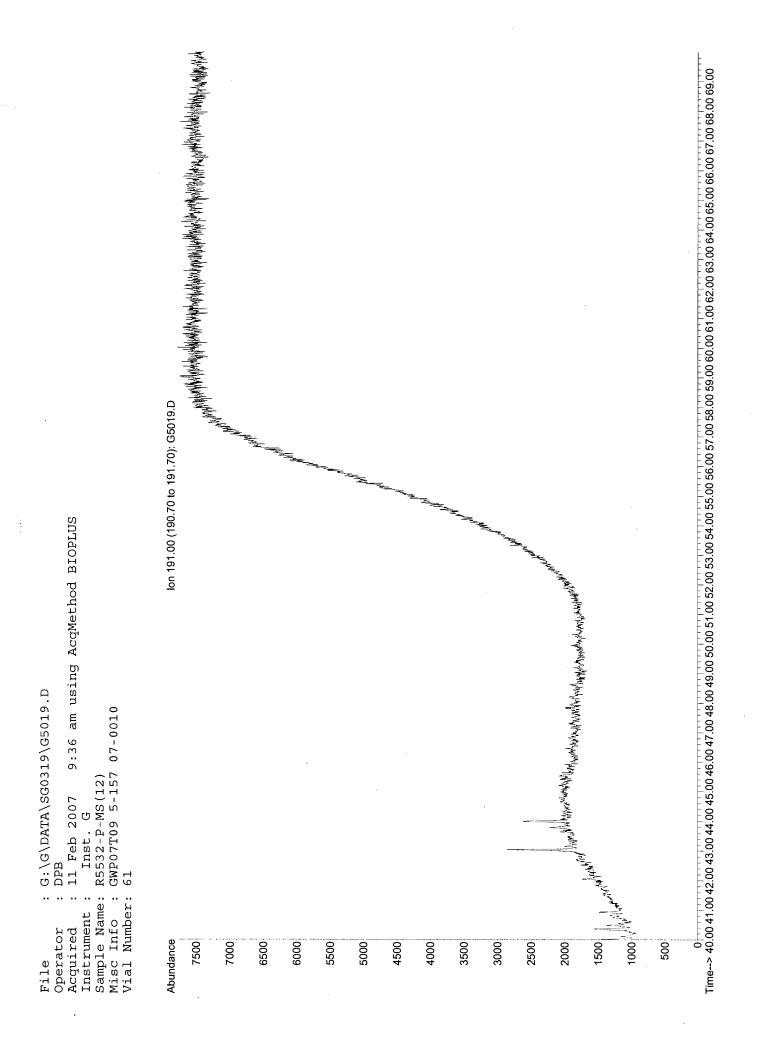


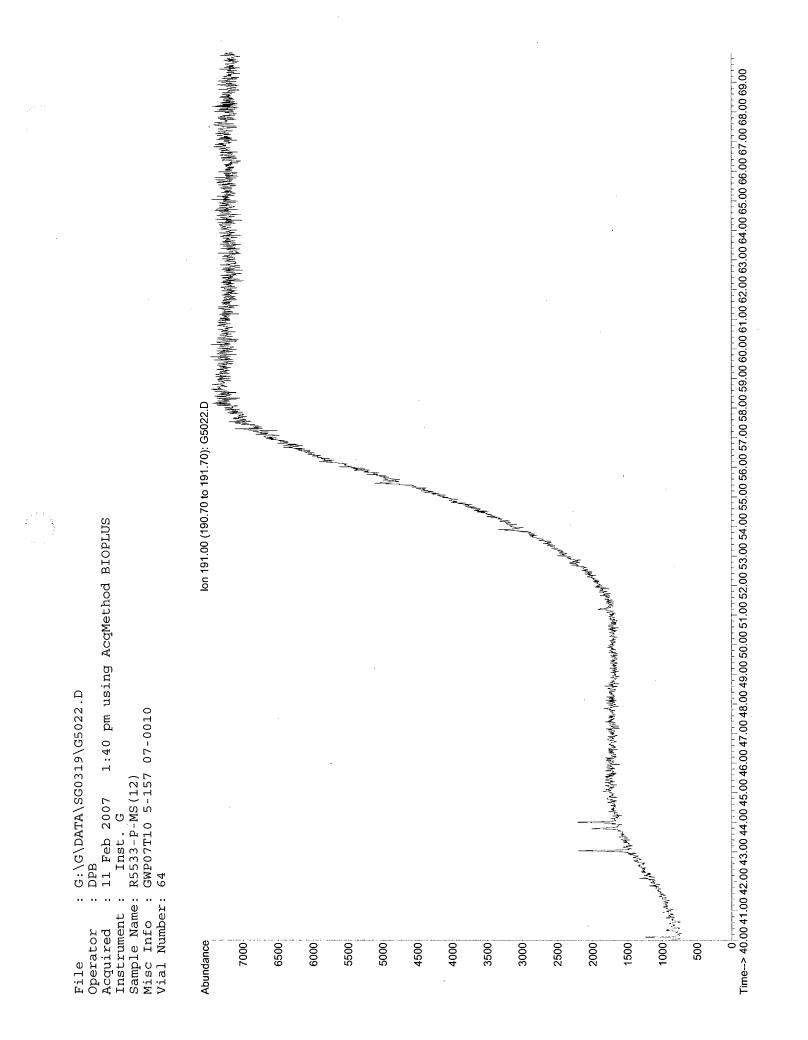


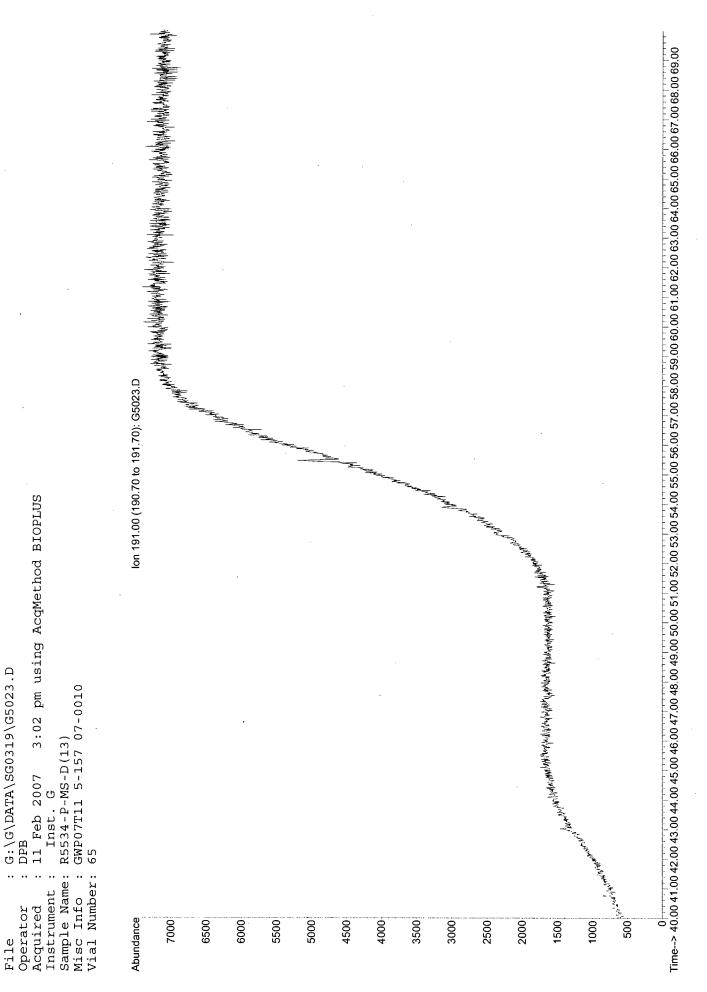


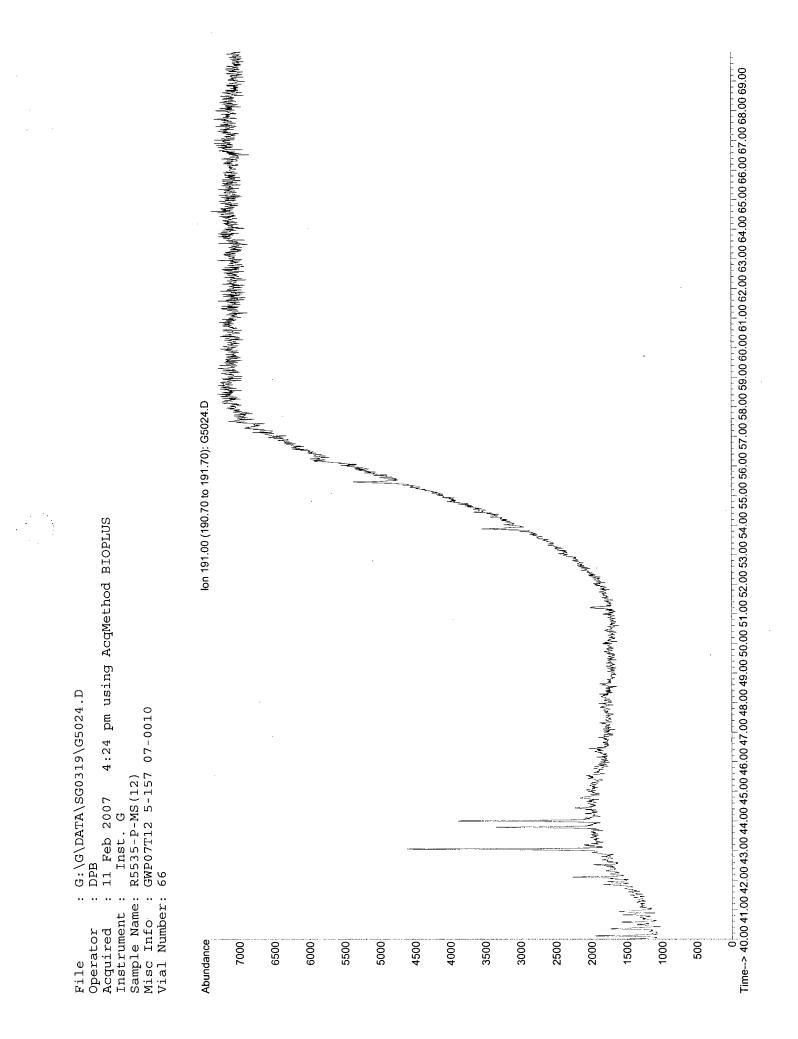


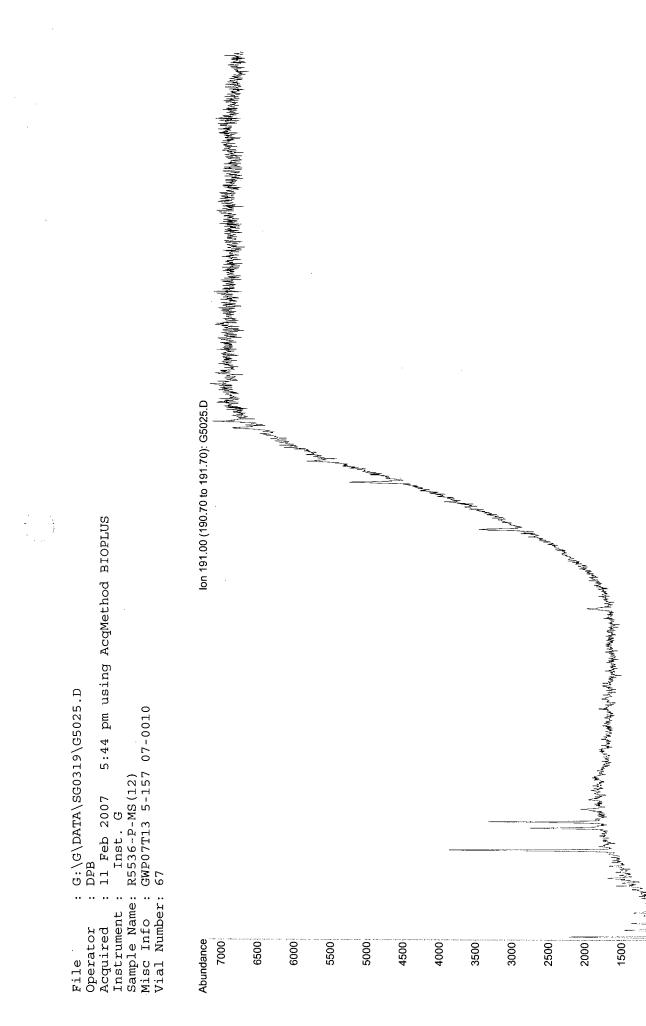
Time--> 40.00 41.00 42.00 43.00 45.00 45.00 46.00 47.00 48.00 49.00 50.00 51.00 52.00 53.00 54.00 55.00 56.00 57.00 58.00 59.00 60.00 61.00 62.00 65.00 66.00 67.00 68.00 69.00







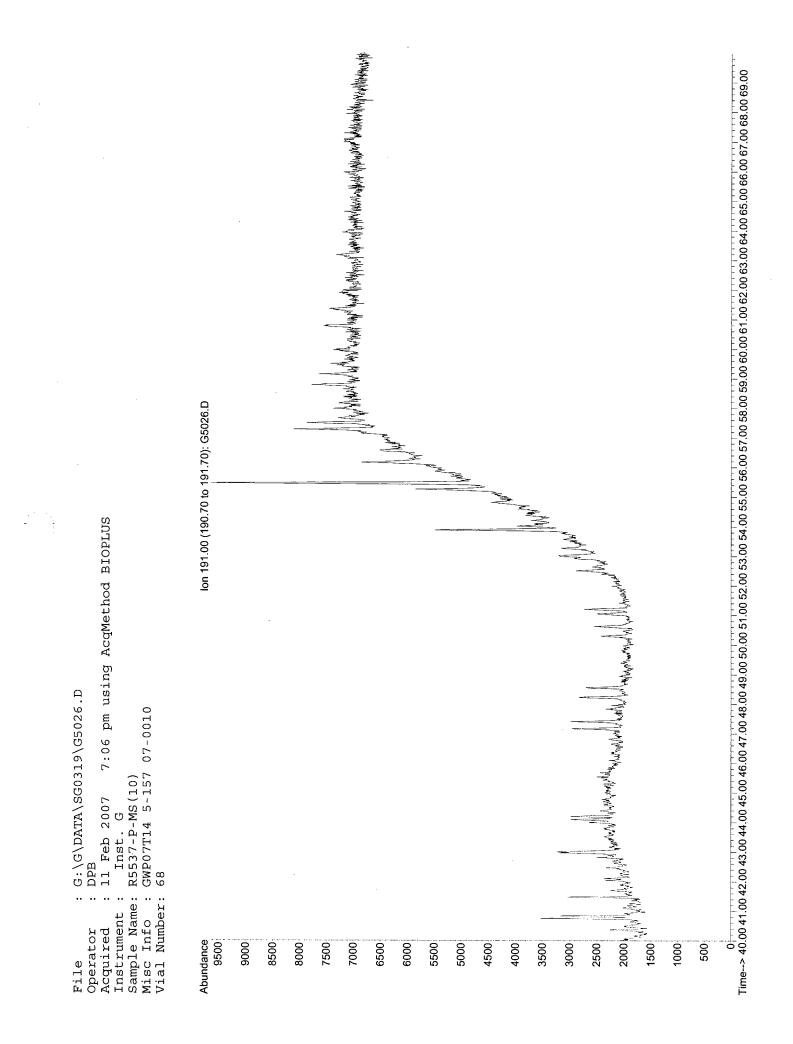


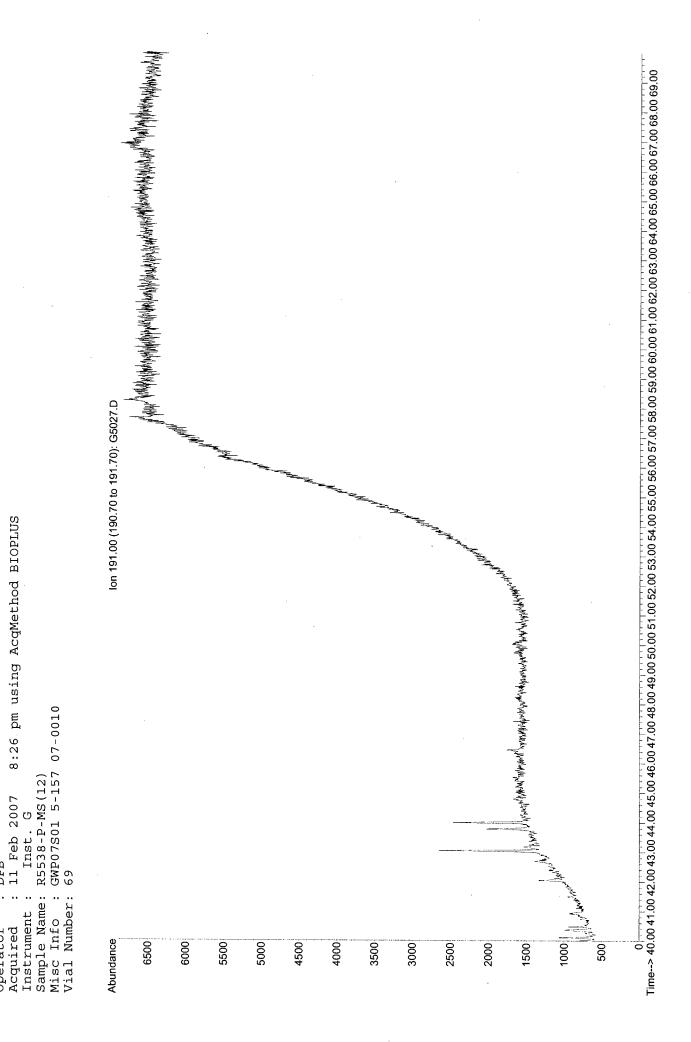


Time--> 40.00 41.00 42.00 45.00 45.00 45.00 45.00 48.00 49.00 50.00 51.00 52.00 53.00 54.00 55.00 56.00 57.00 58.00 59.00 60.00 61.00 62.00 65.00 65.00 68.00 69.00 Ö

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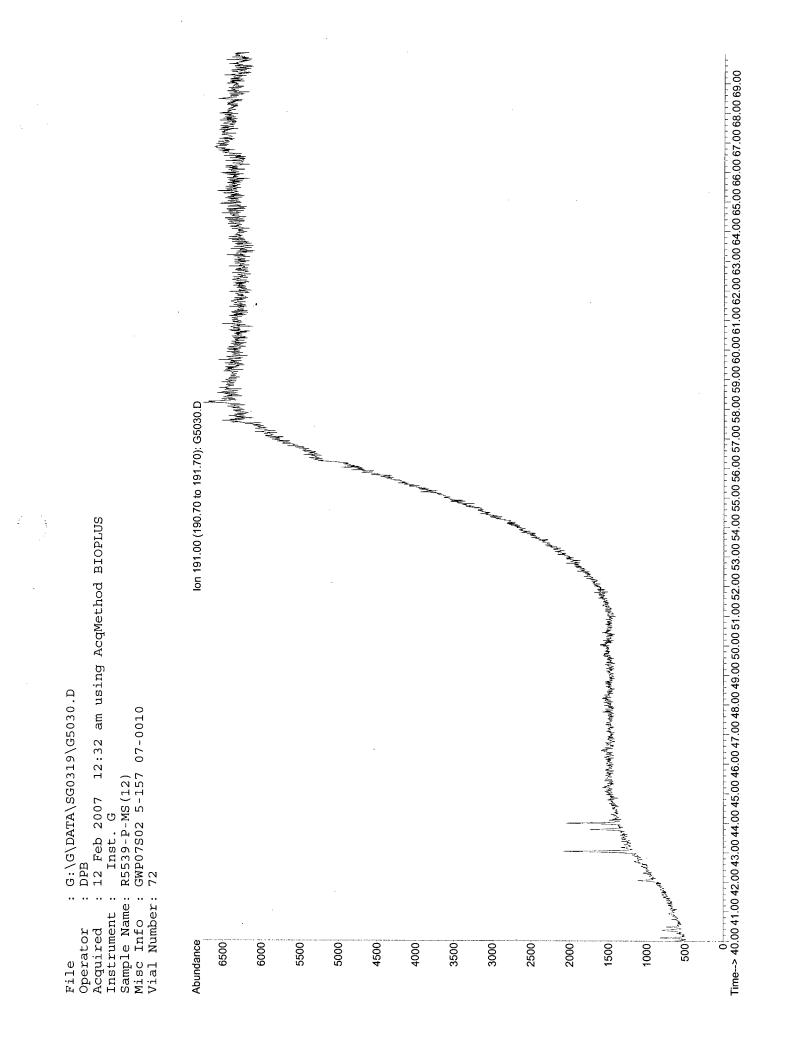


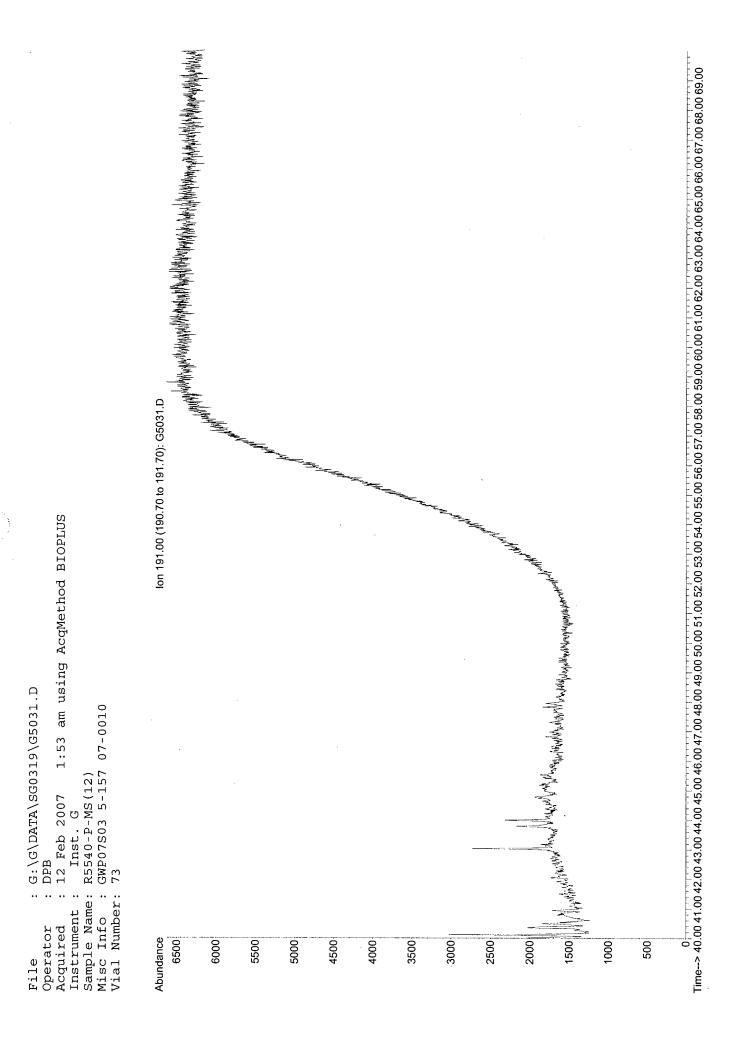


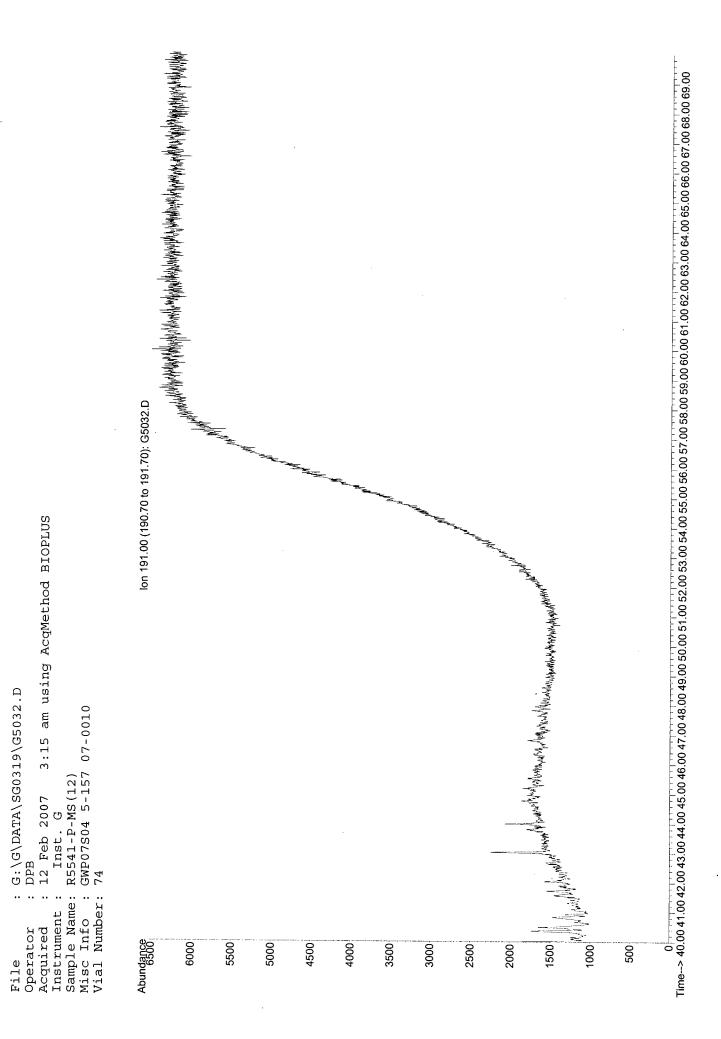
G:\G\DATA\SG0319\G5027.D DPB

Operator

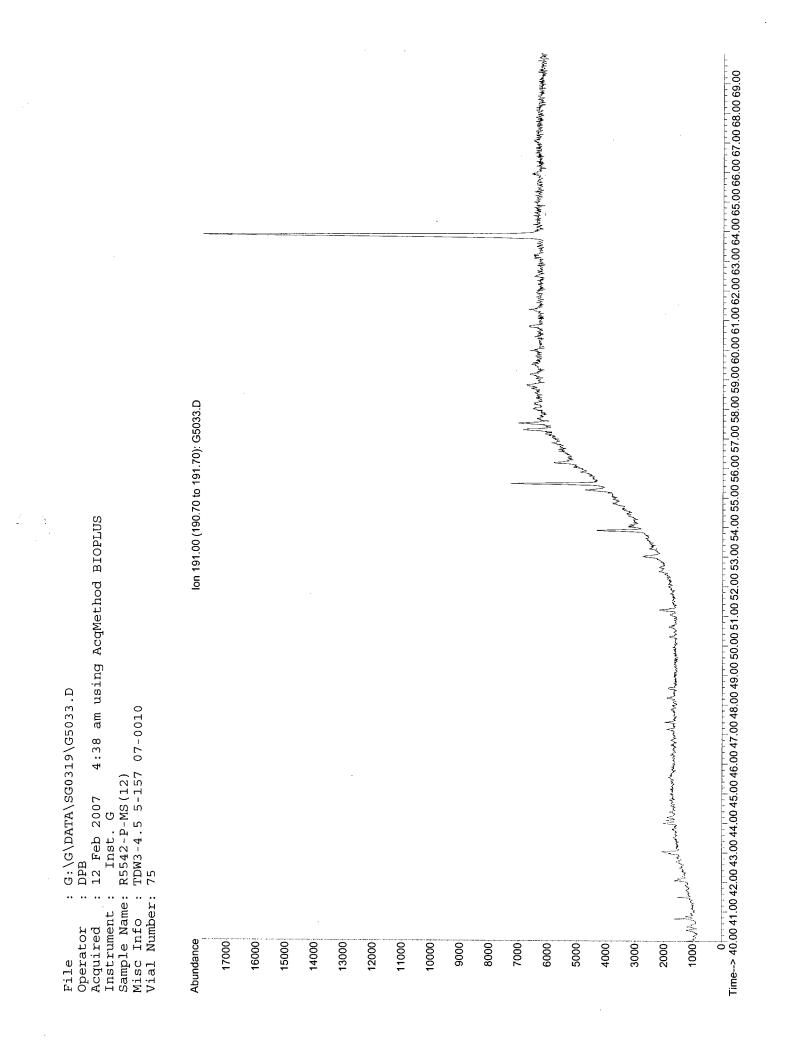
File

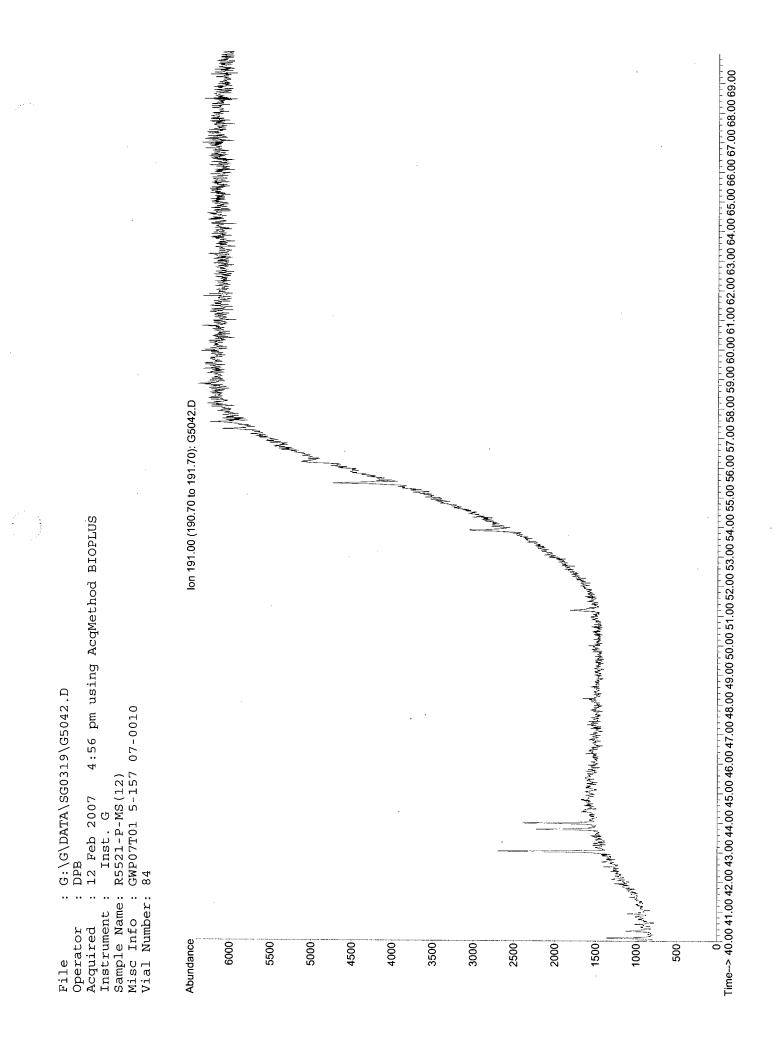


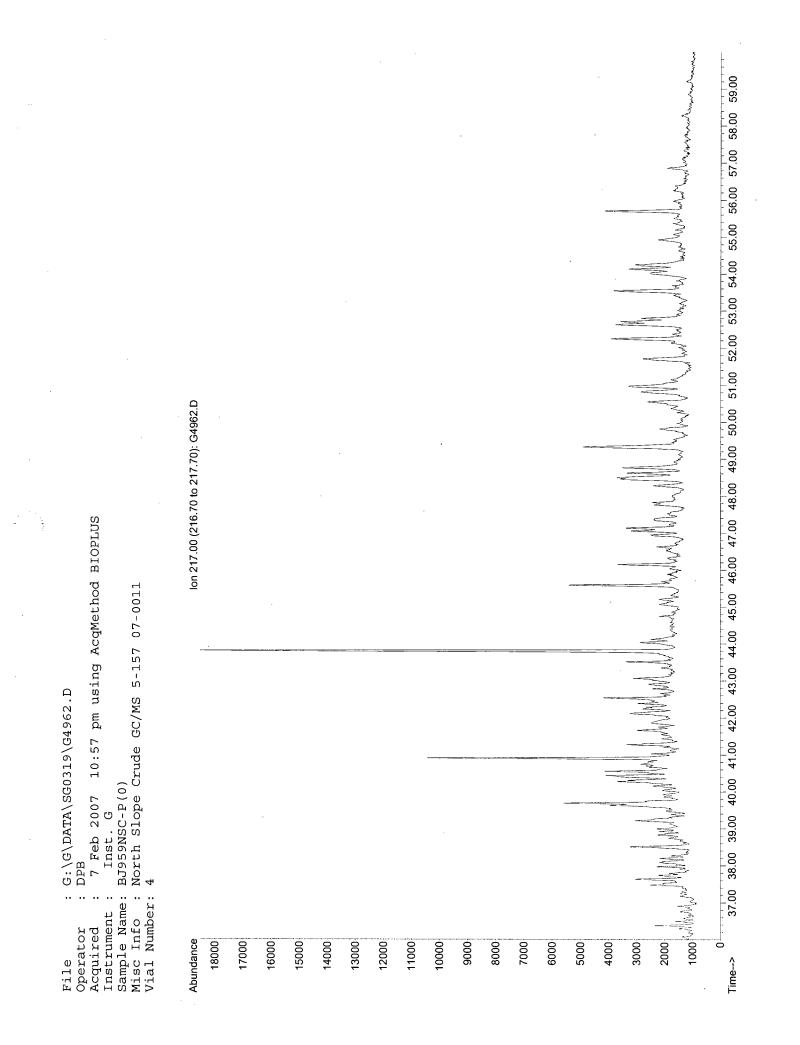


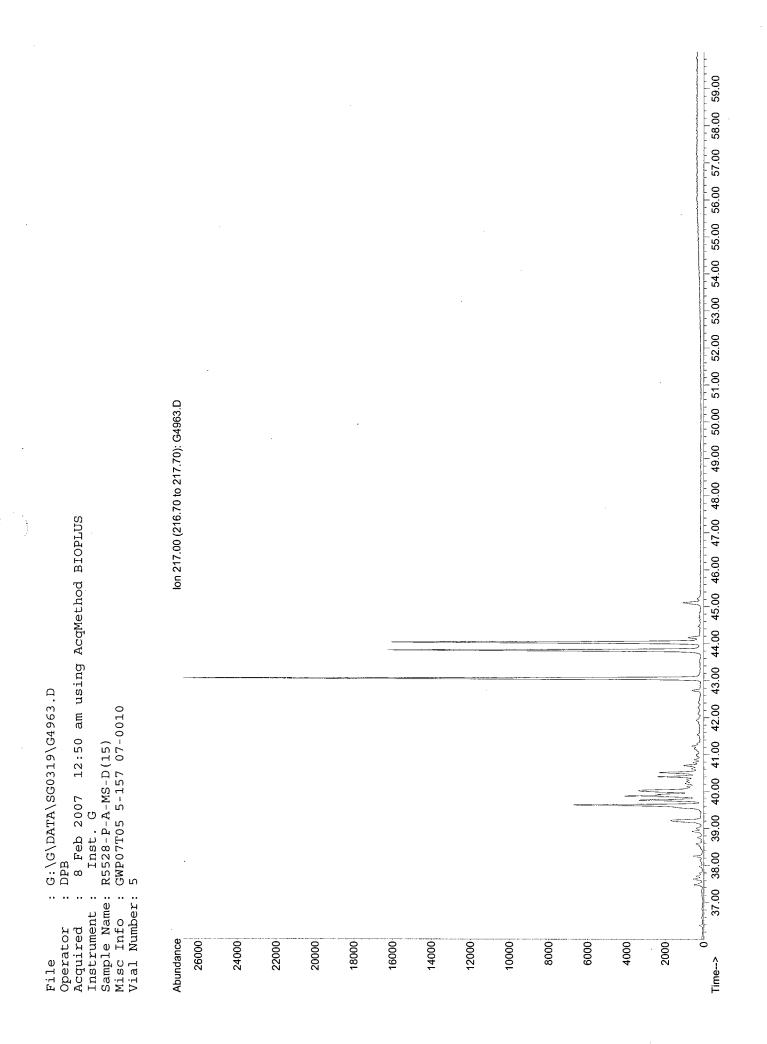


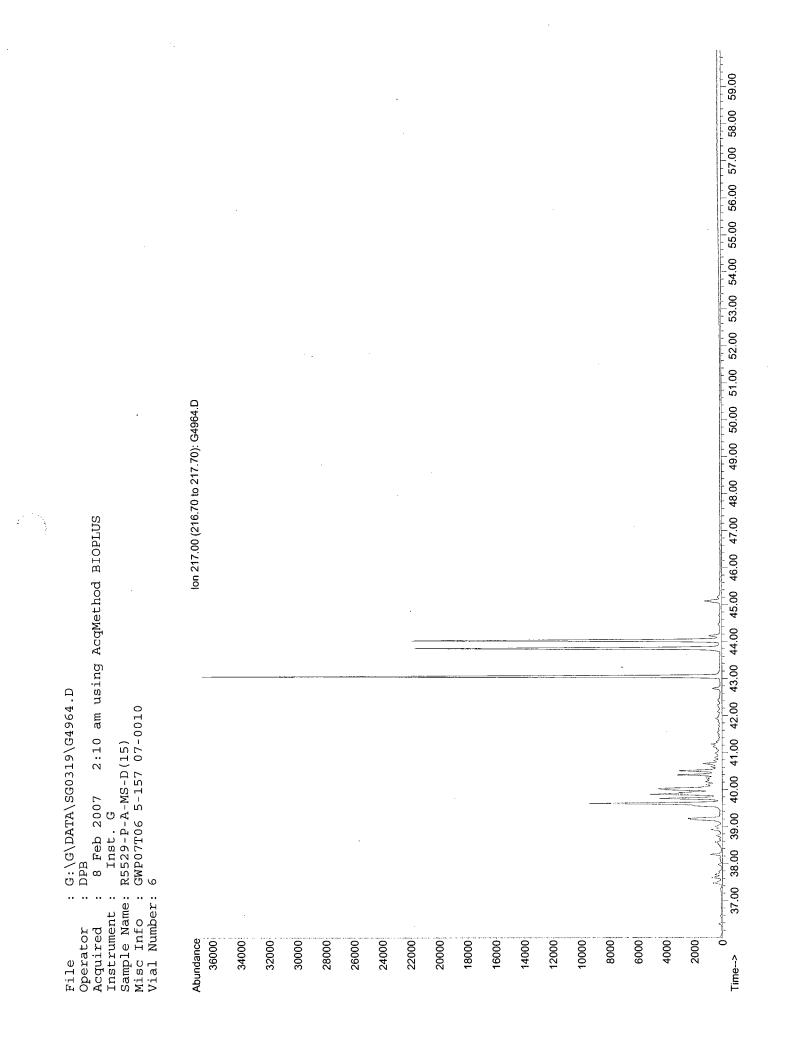
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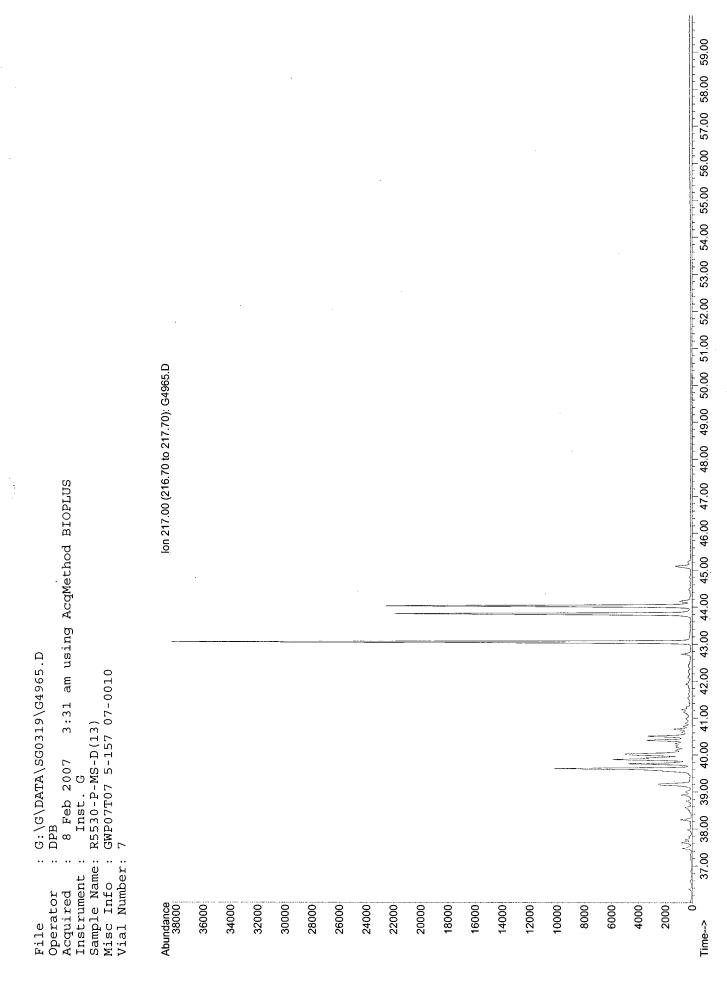




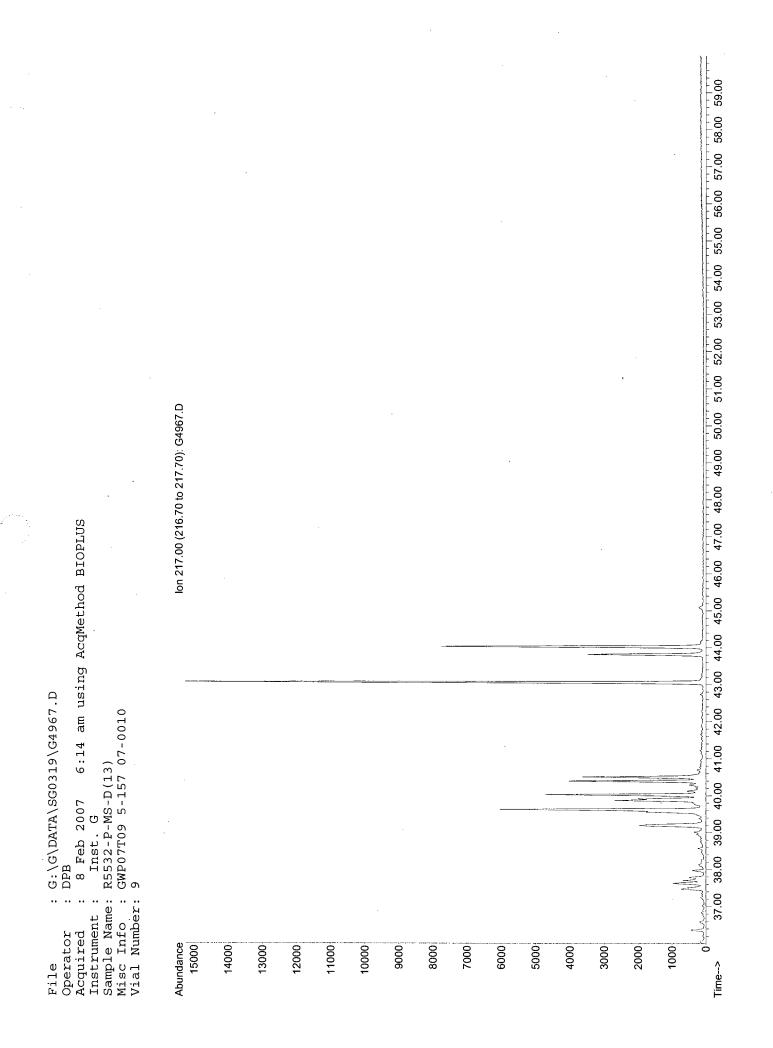


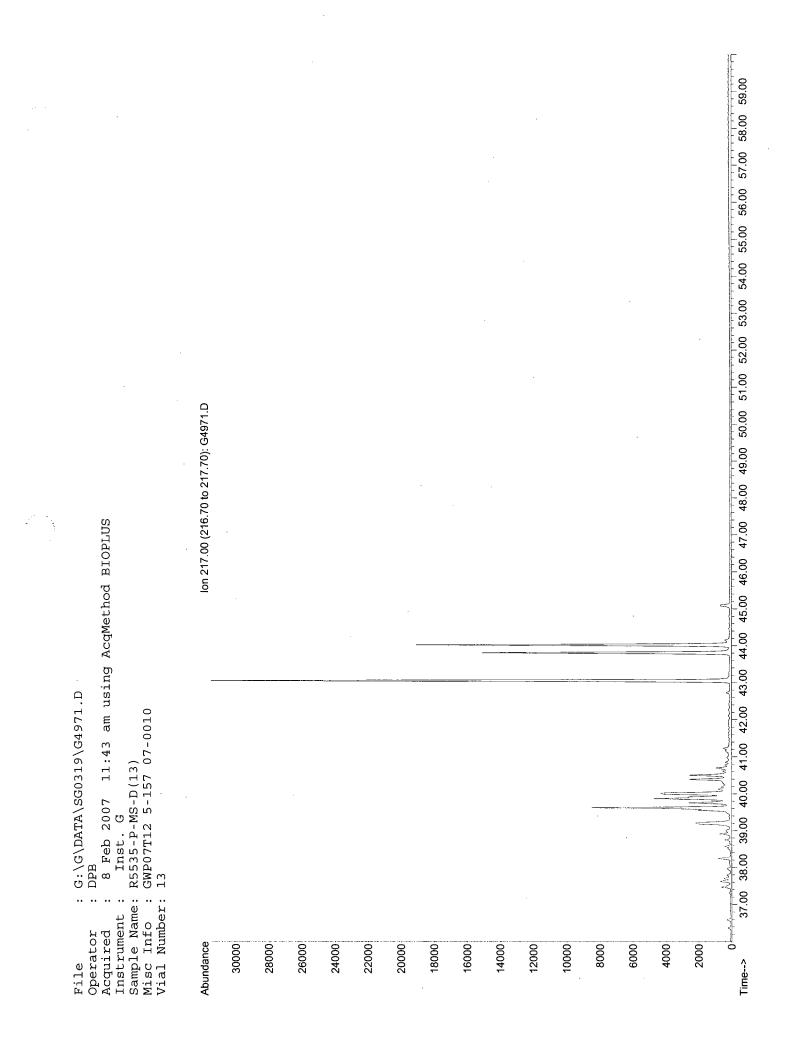


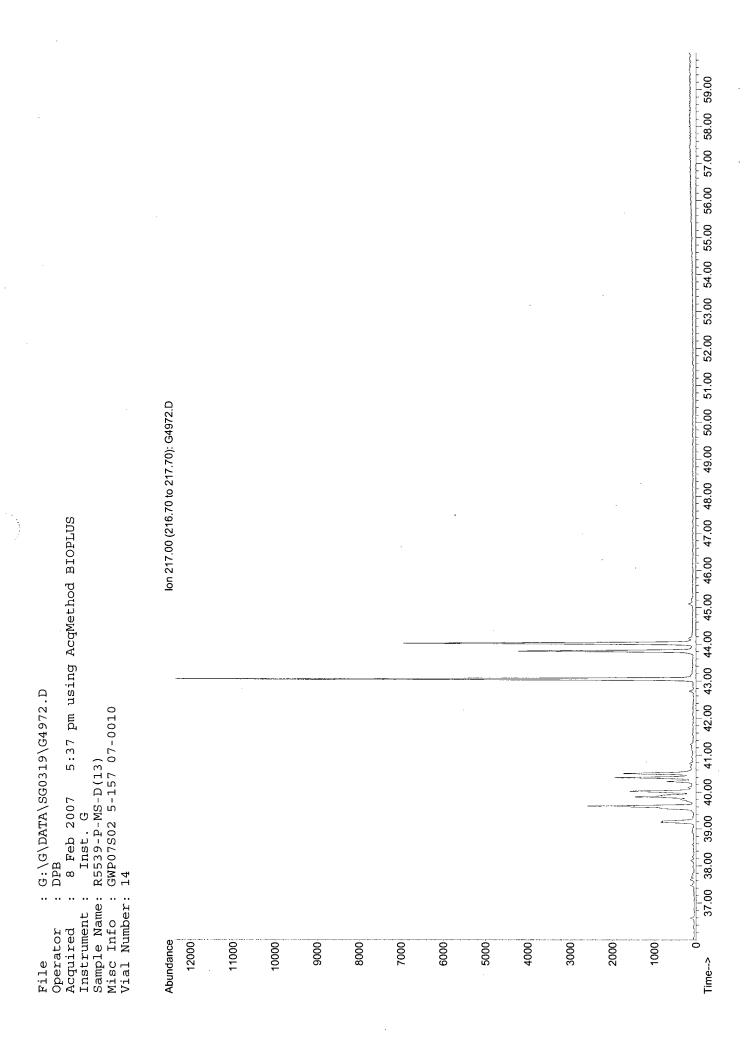


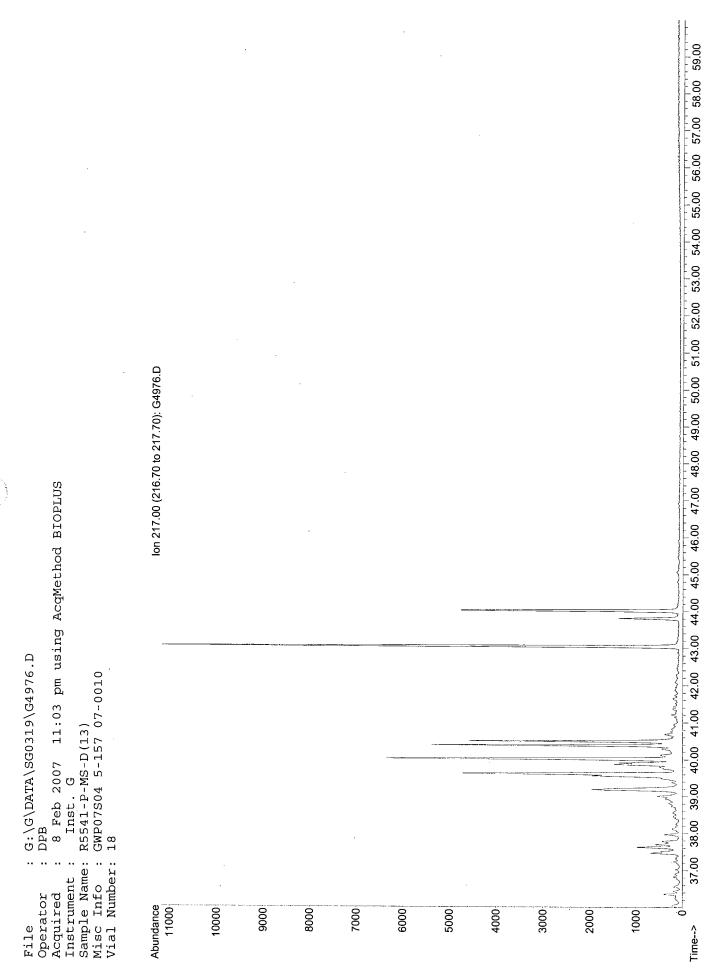


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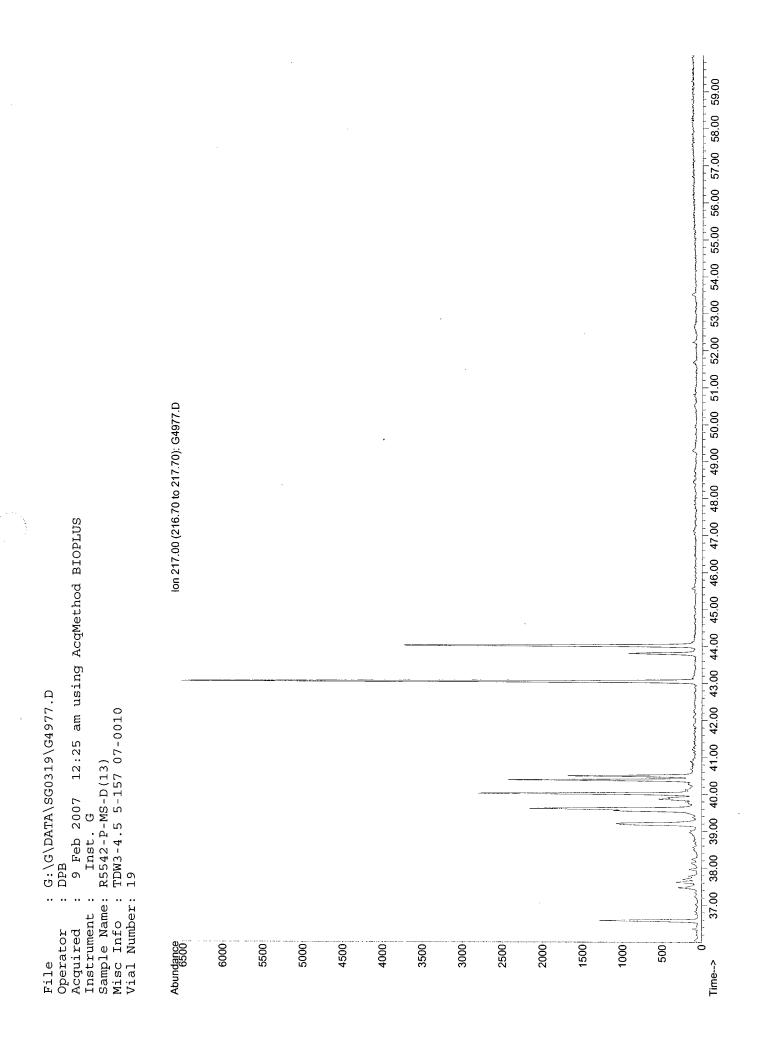


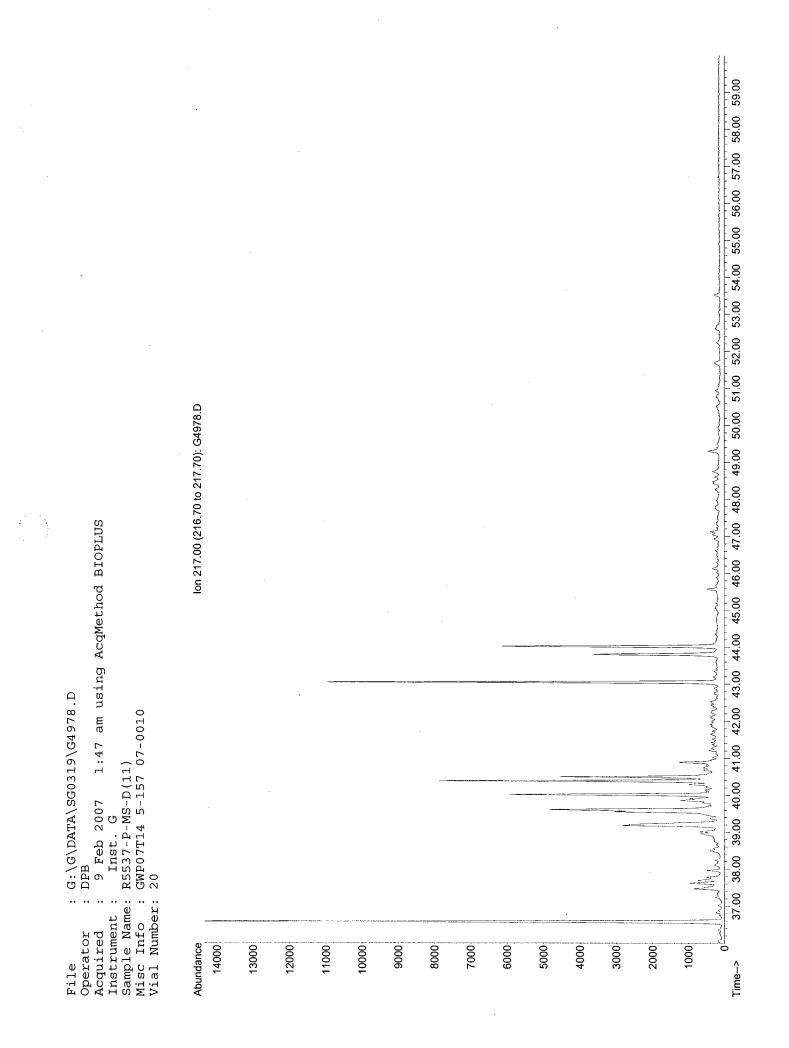


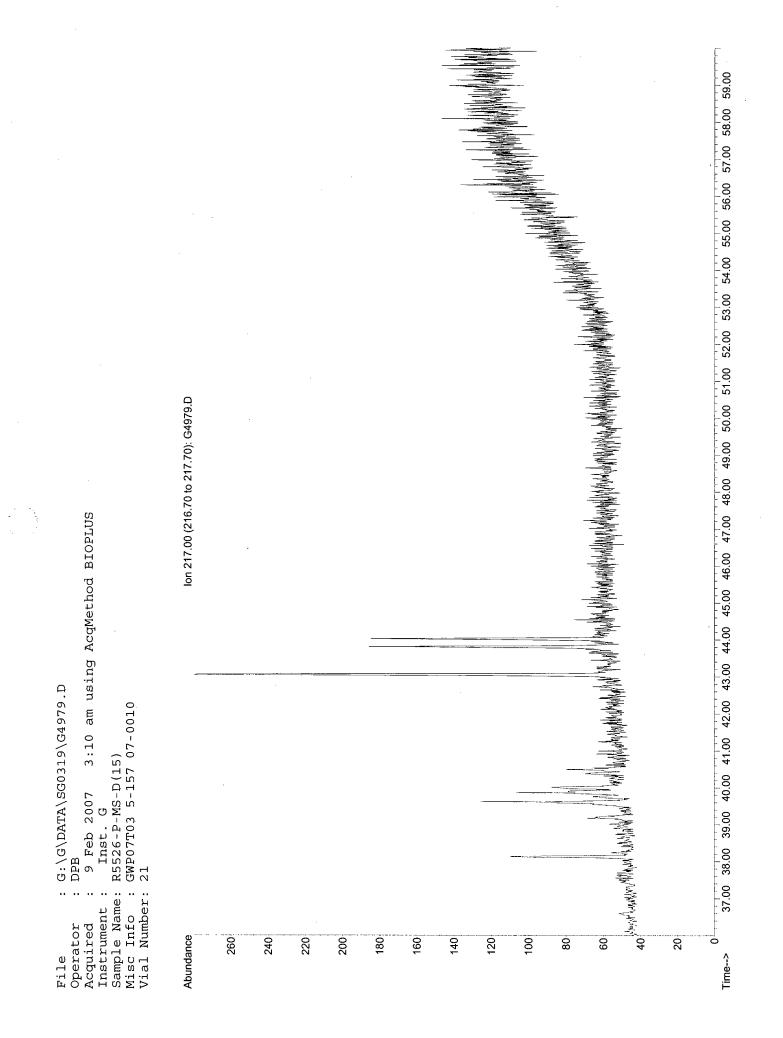


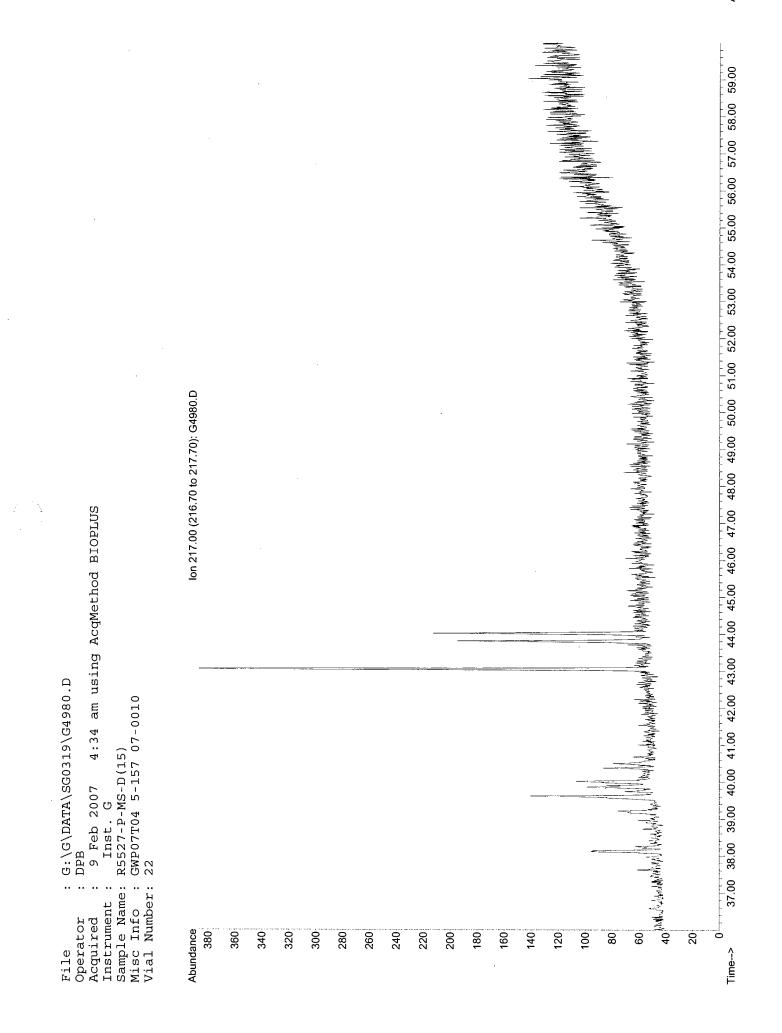


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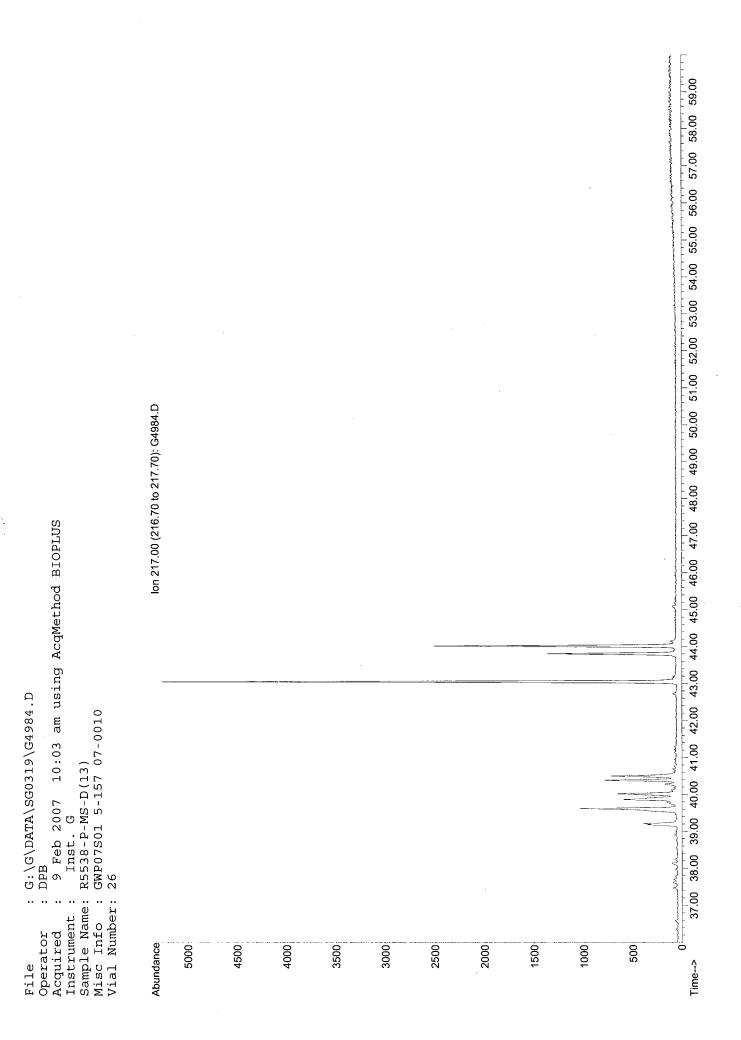


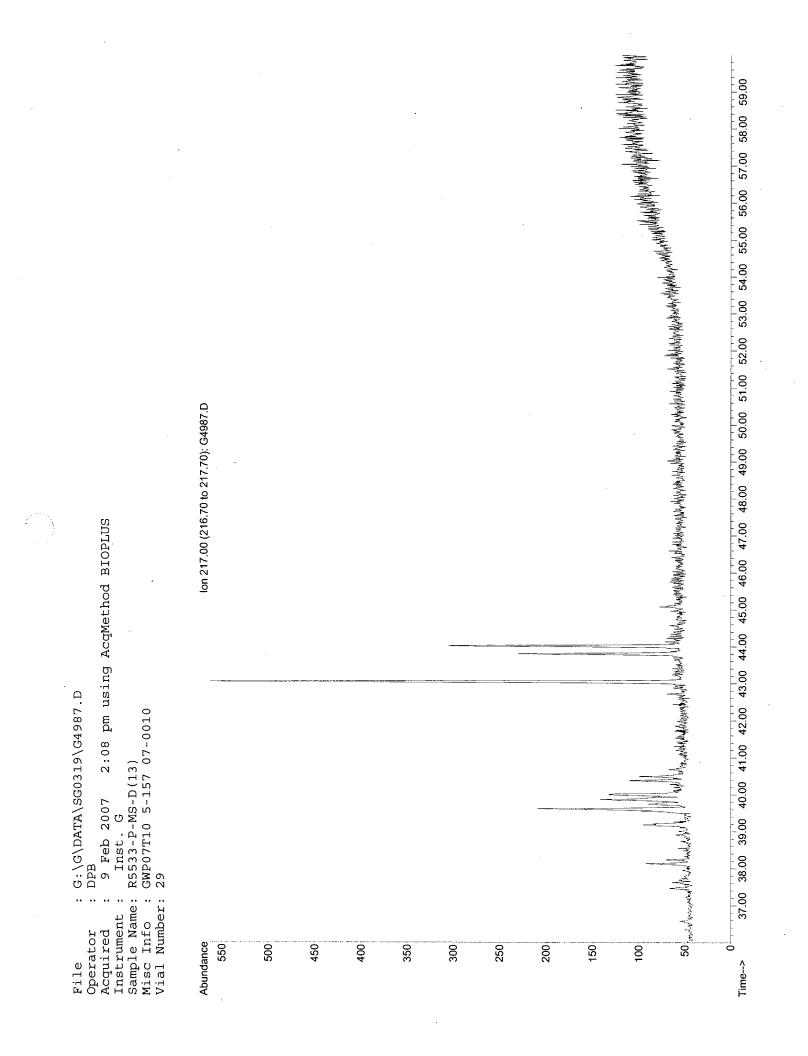
37.00 38.00 39.00 40.00 41.00 42.00 43.00 44.00 45.00 46.00 47.00 48.00 49.00 50.00 51.00 52.00 54.00 55.00 56.00 57.00 58.00 59.00 Ion 217.00 (216.70 to 217.70); G4983.D 8:41 am using AcqMethod BIOPLUS : G:\G\DATA\SG0319\G4983.D : DPB Auguired : 9 Feb 2007 8:41 am
Instrument : Inst.G
Sample Name: R5536-P-MS-D(13)
Misc Info : GWP07T13 5-157 07-0010
Vial Number: 25 Niv. Abundance 12000 11000 10000 0006 8000 7000 6000 5000 4000 3000 1000 2000

Time-->

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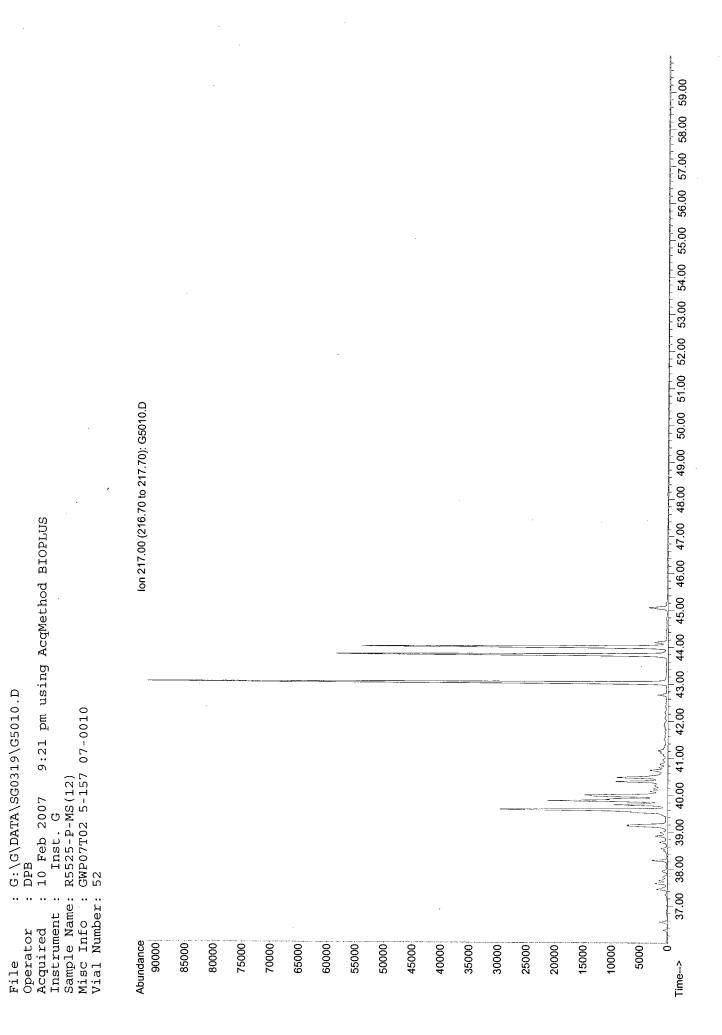
File



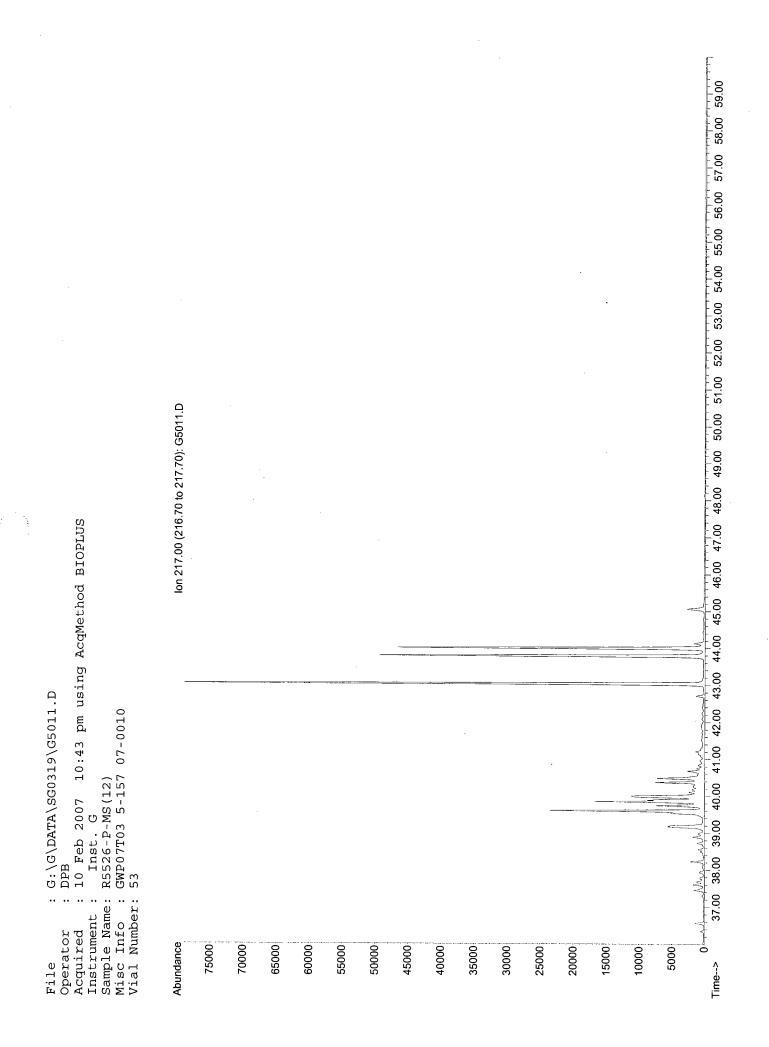


37.00 38.00 39.00 40.00 41.00 42.00 43.00 44.00 45.00 46.00 47.00 48.00 49.00 50.00 51.00 52.00 53.00 54.00 55.00 56.00 57.00 58.00 59.00 lon 217.00 (216.70 to 217.70): G5007.D 5:18 pm using AcqMethod BIOPLUS Acquired : 10 Feb 2007 5:18 pm using Act Instrument : Inst. G Sample Name: BJ939PB-P-MS(5) Misc Info : Procedural Blank 5-157 07-0010 Vial Number: 49 G:\G\DATA\SG0319\G5007.D DPB ... Abundance 120000 110000 00006 70000 60000 100000 80000 50000 40000 30000 20000 10000 0 File Time-->

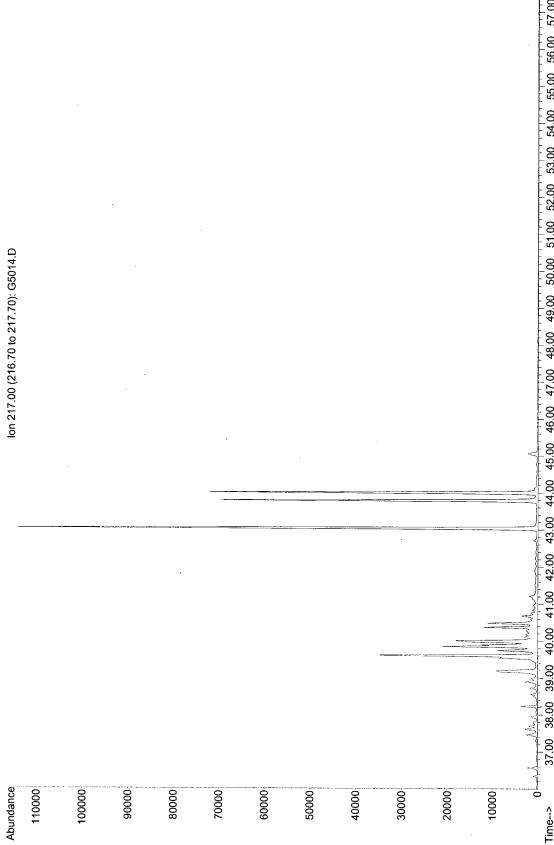
37.00 38.00 39.00 40.00 41.00 42.00 43.00 44.00 45.00 46.00 47.00 48.00 49.00 50.00 51.00 52.00 53.00 54.00 55.00 56.00 57.00 58.00 59.00 lon 217.00 (216.70 to 217.70); G5008.D 6:37 pm using AcqMethod BIOPLUS Laboratory Control Sample 5-157 07-0010 G:\G\DATA\SG0319\G5008.D DPB Sample Name: BJ940LCS-P-MS(5) Misc Info : Laboratory Contr Vial Number: 50 10 Feb 2007 Inst. G •• •• Instrument Acquired Operator Abundance 120000 110000 100000 00006 80000 70000 60000 50000 40000 30000 20000 10000 ò File Time-->



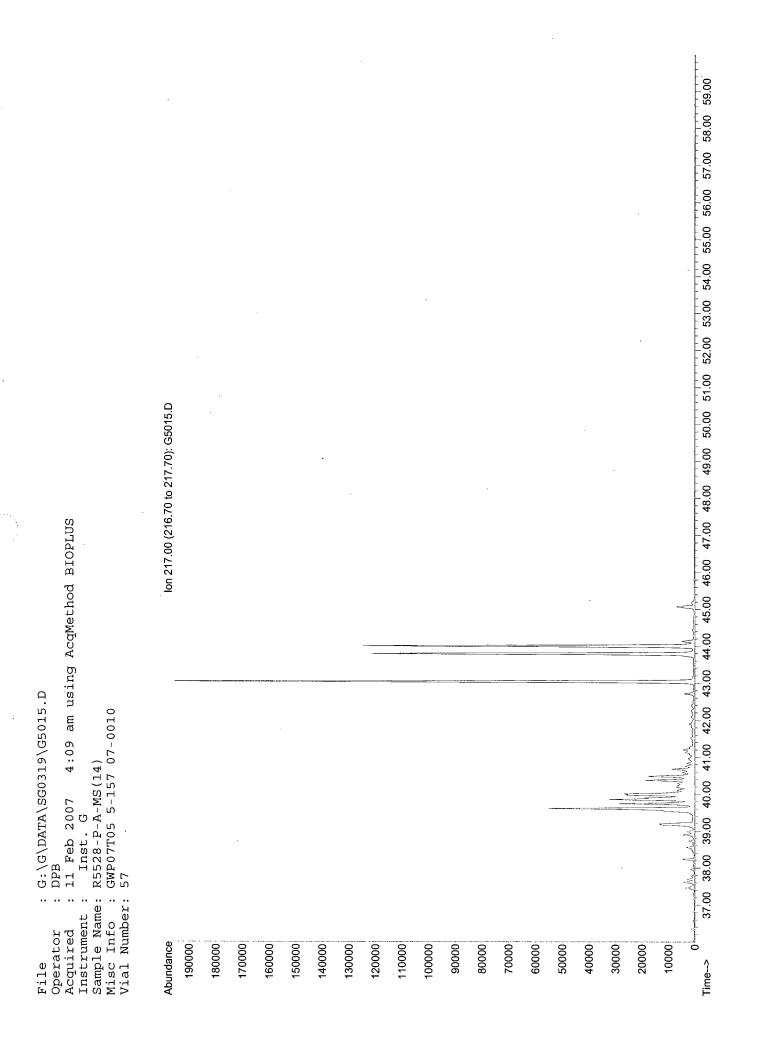
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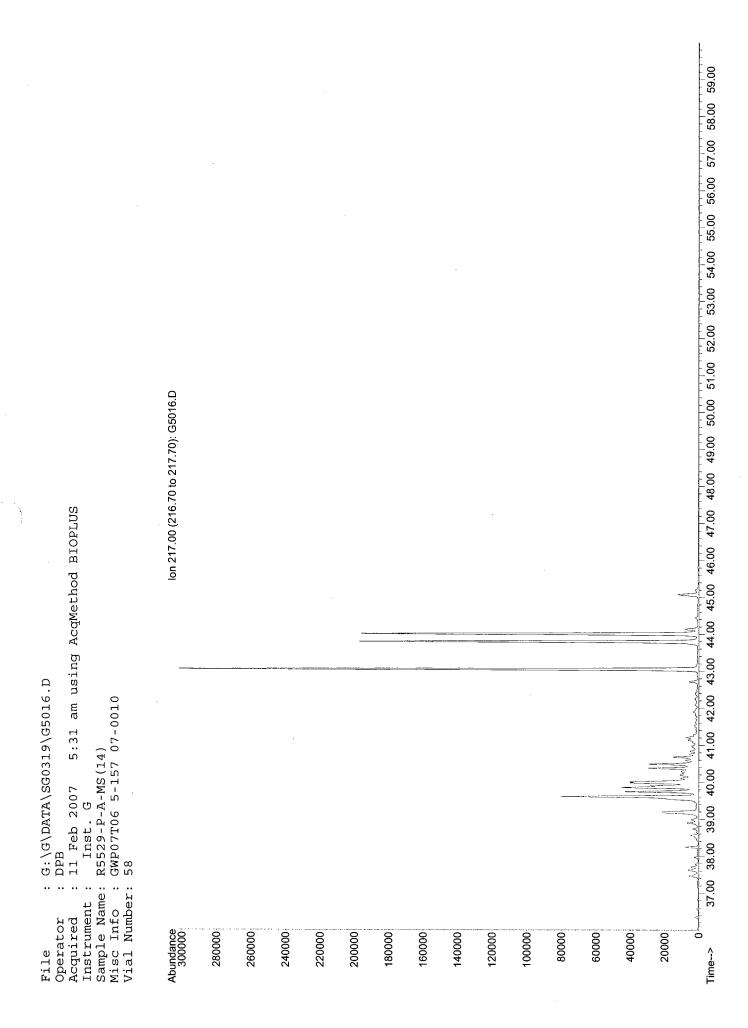


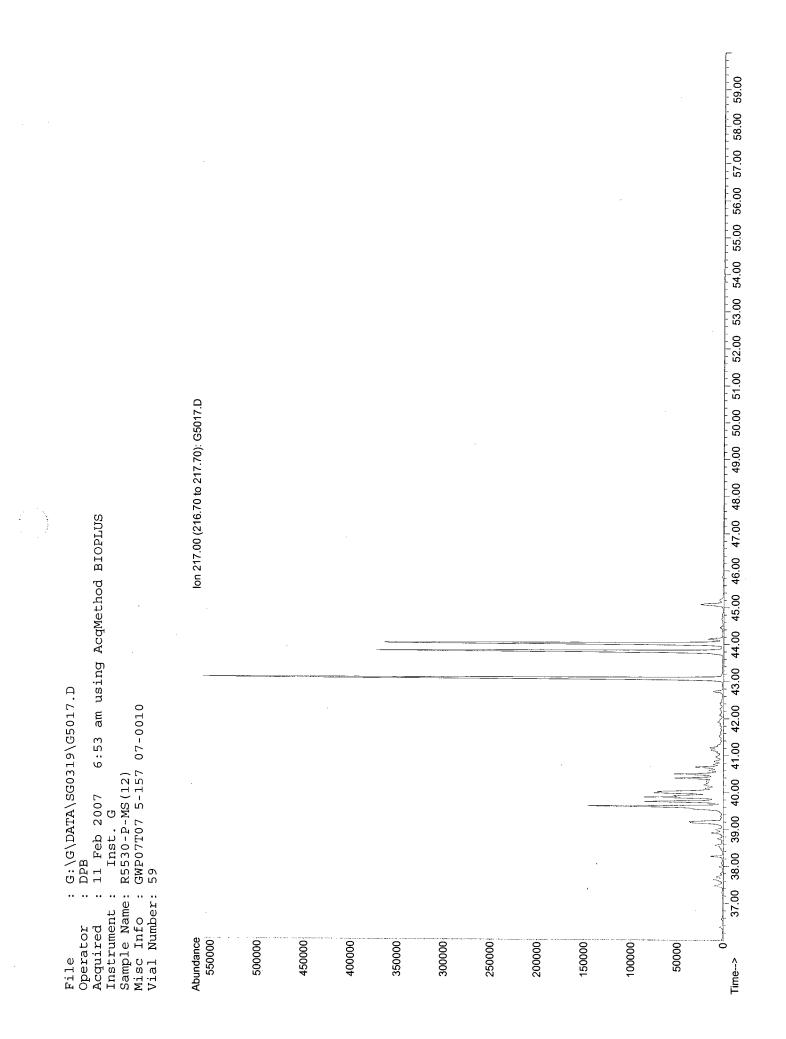
lon 217.00 (216.70 to 217.70): G5014.D 2:47 am using AcqMethod BIOPLUS G:\G\DATA\SG0319\G5014.D DPB Acquired : 11 Feb 2007 2:47 am Instrument : Inst. G Sample Name: R5527-P-MS(12) Misc Info : GWP07T04 5-157 07-0010 Vial Number: 56 Operator File

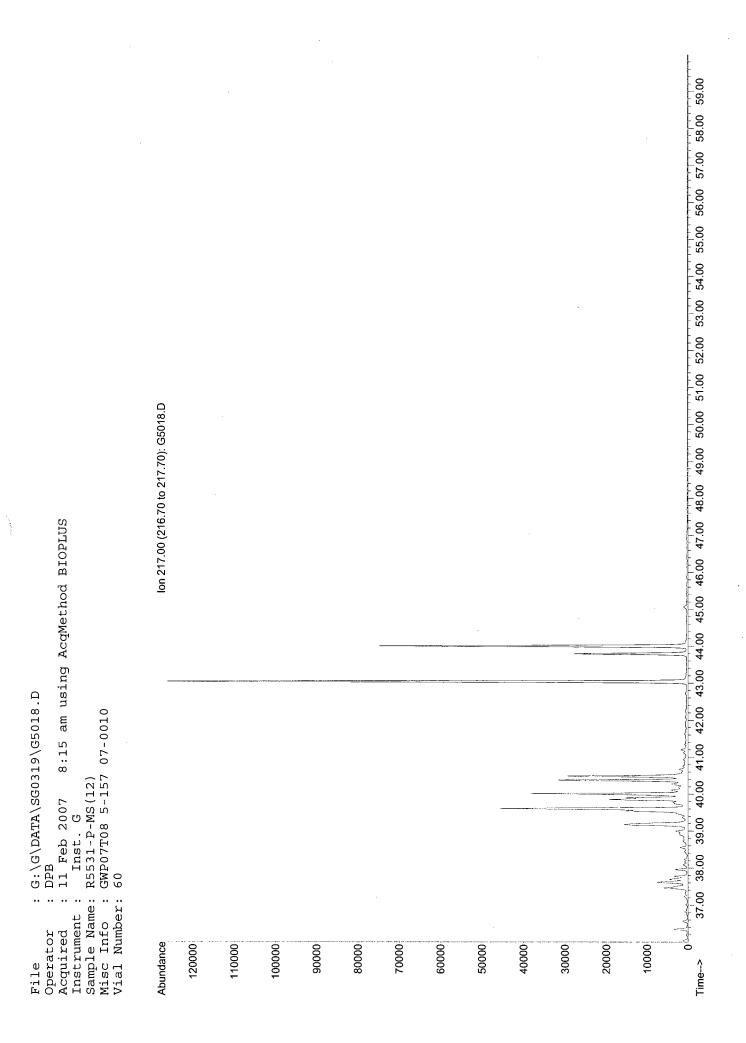


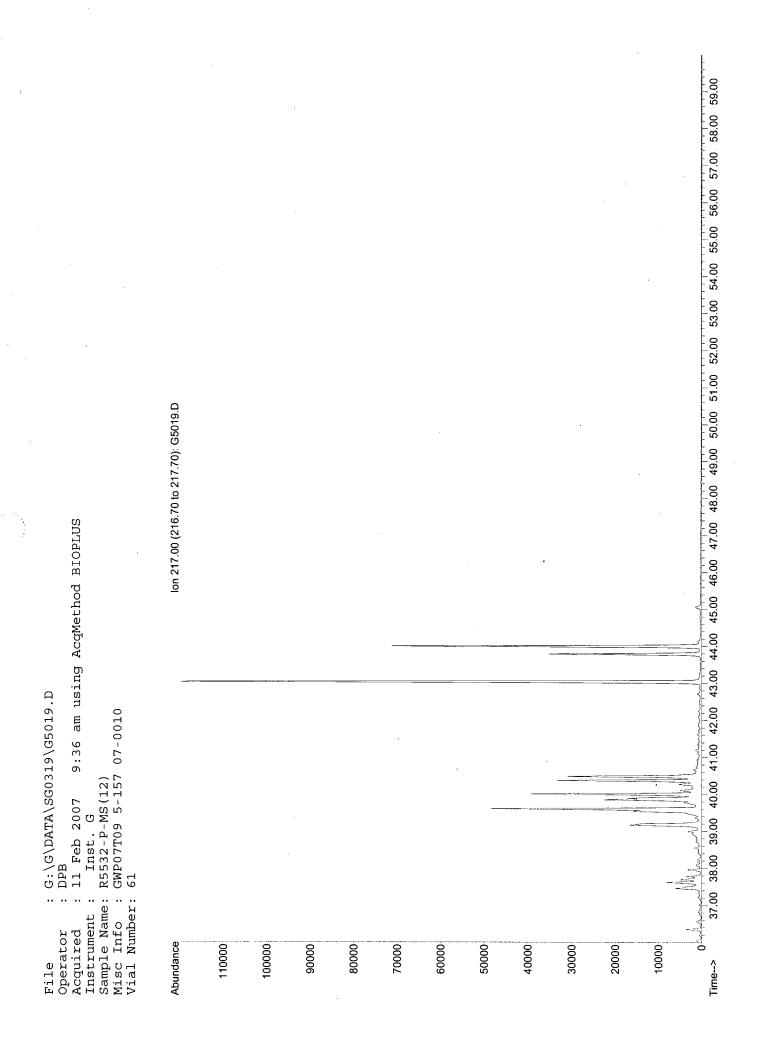
37.00 38.00 39.00 40.00 41.00 42.00 44.00 45.00 45.00 47.00 48.00 49.00 50.00 51.00 52.00 53.00 55.00 56.00 57.00 58.00 59.00



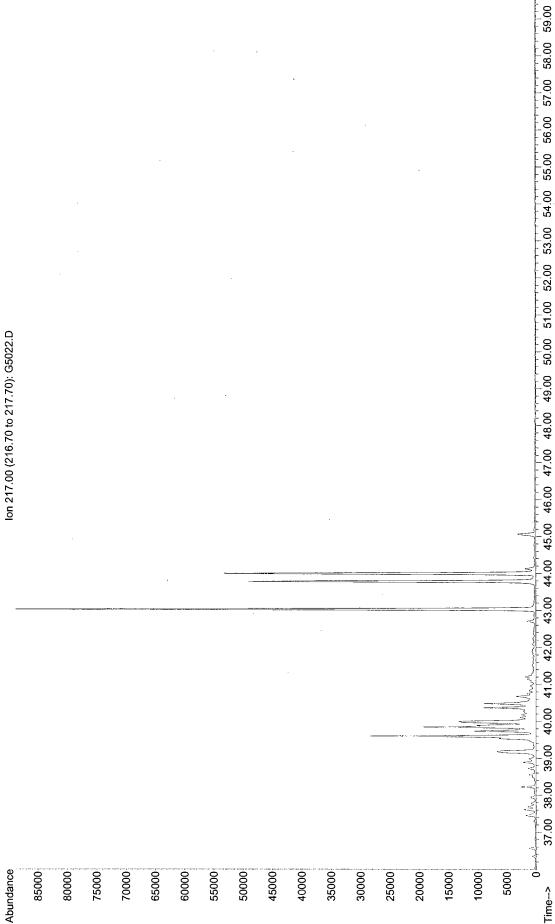




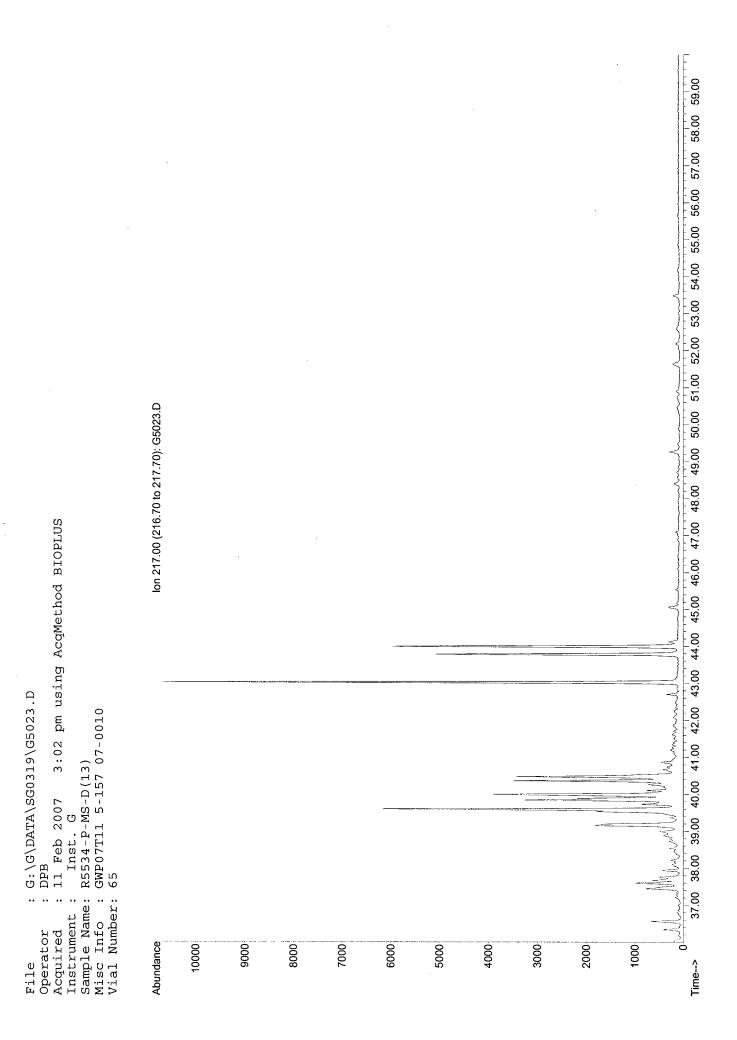


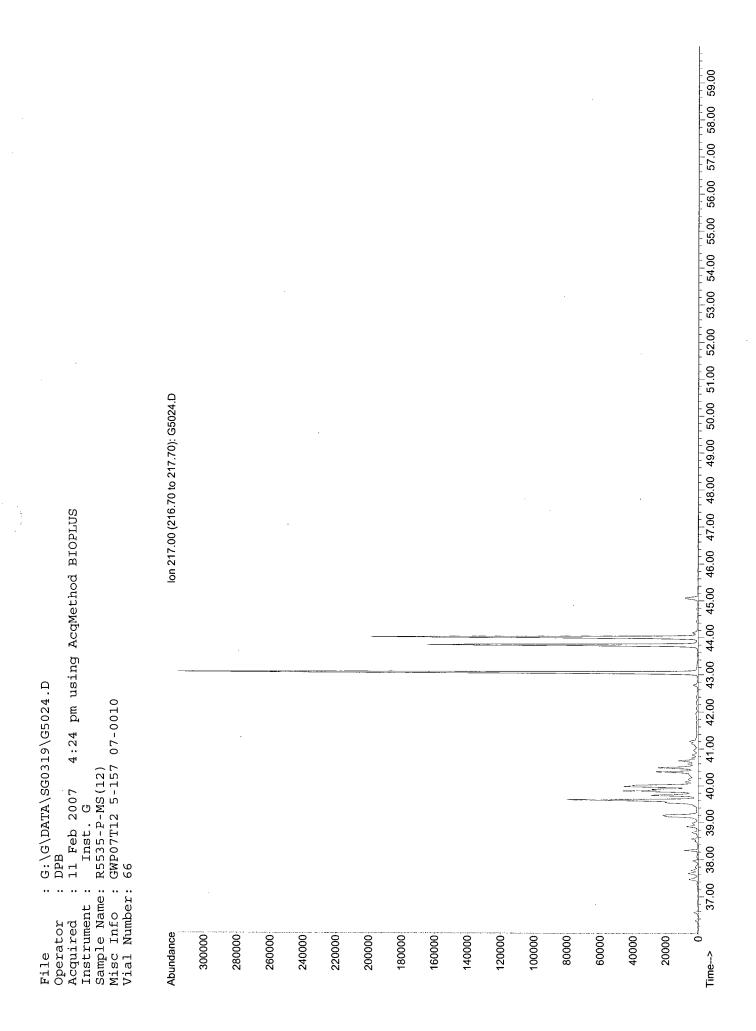


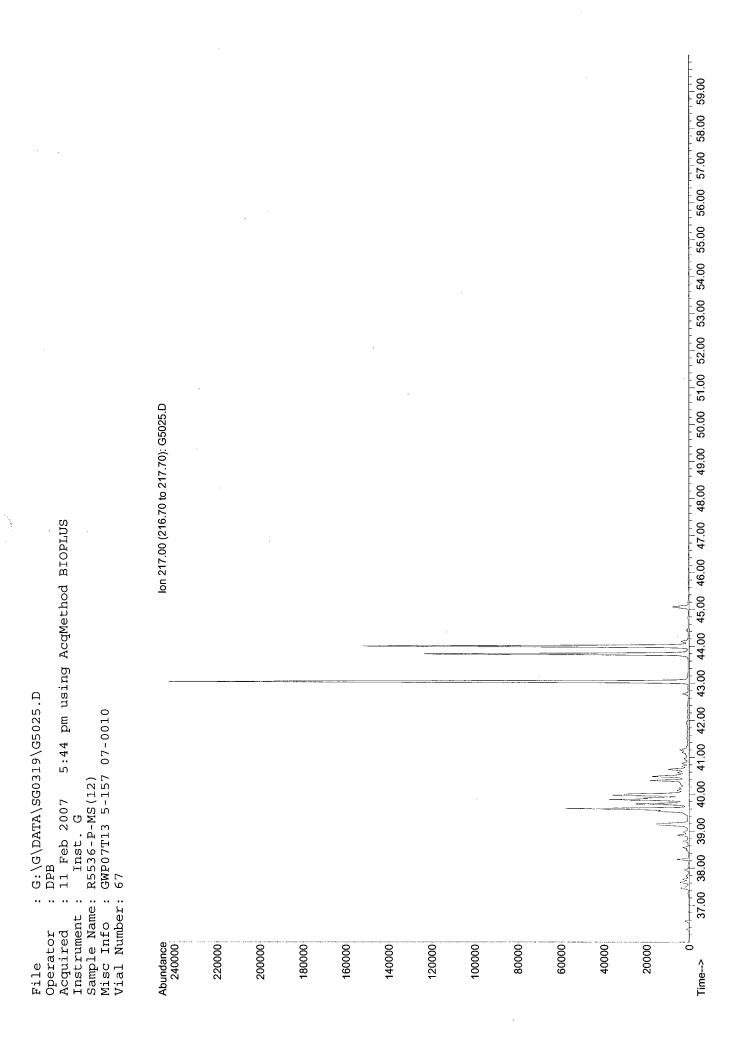
1:40 pm using AcqMethod BIOPLUS G:\G\DATA\SG0319\G5022.D DPB Acquired : 11 Feb 2007 1:40 pm 1 Instrument : Inst. G Sample Name: R5533-P-MS(12) Misc Info : GWP07T10 5-157 07-0010 Vial Number: 64 Operator File

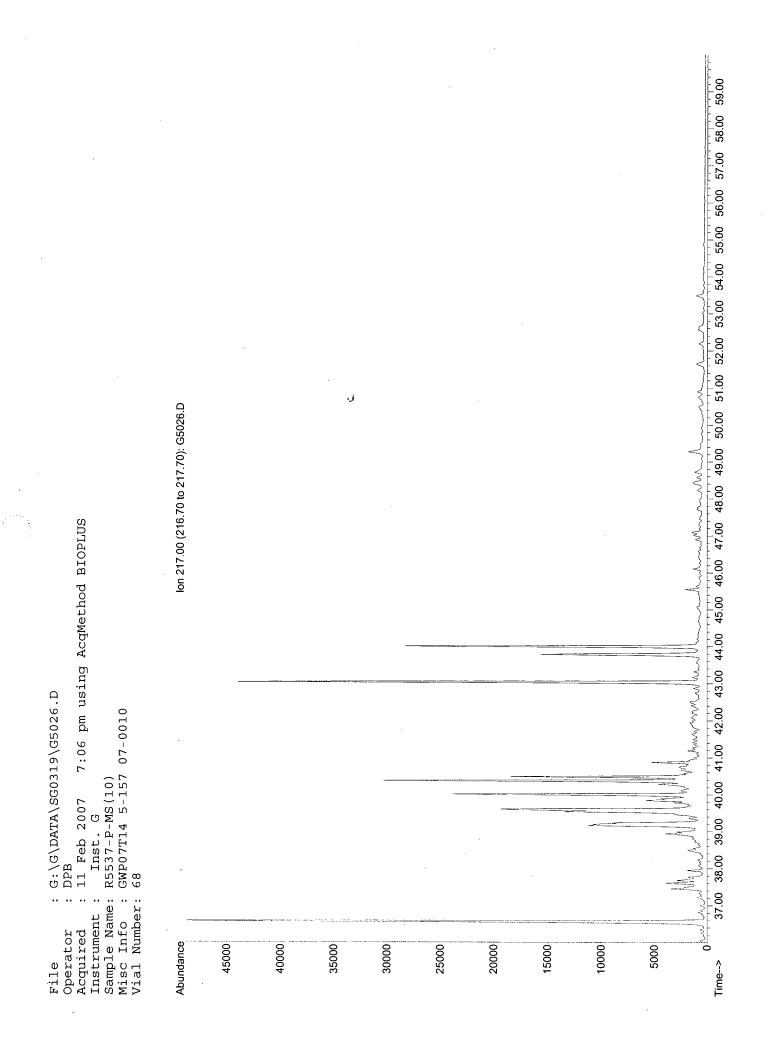


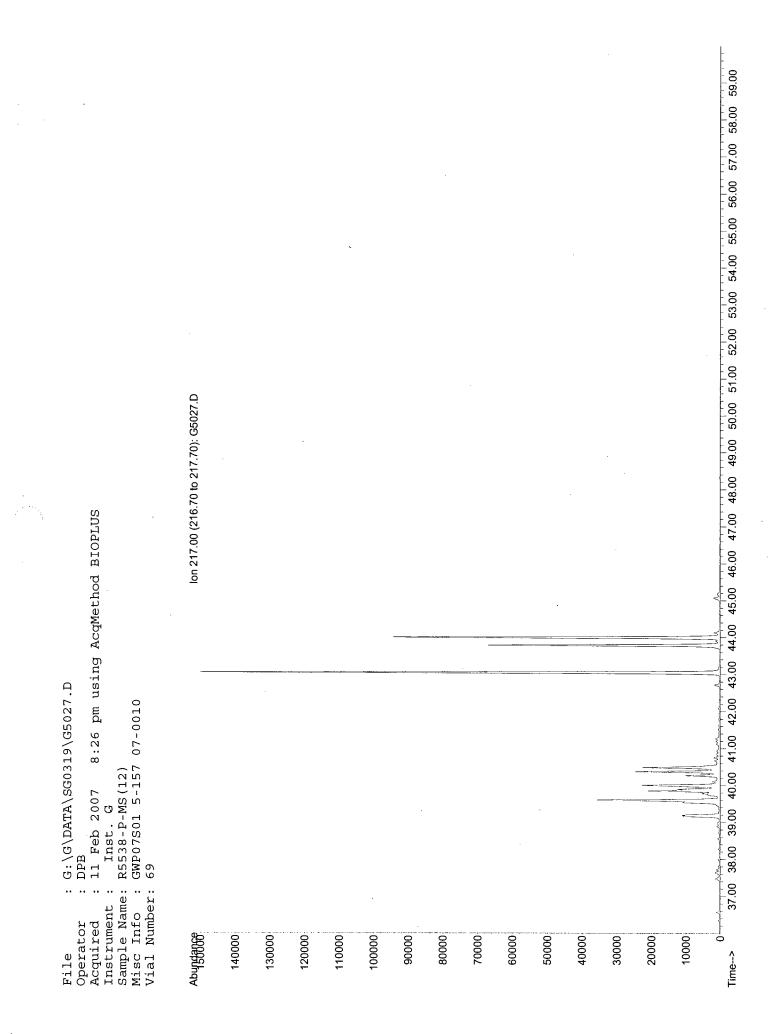
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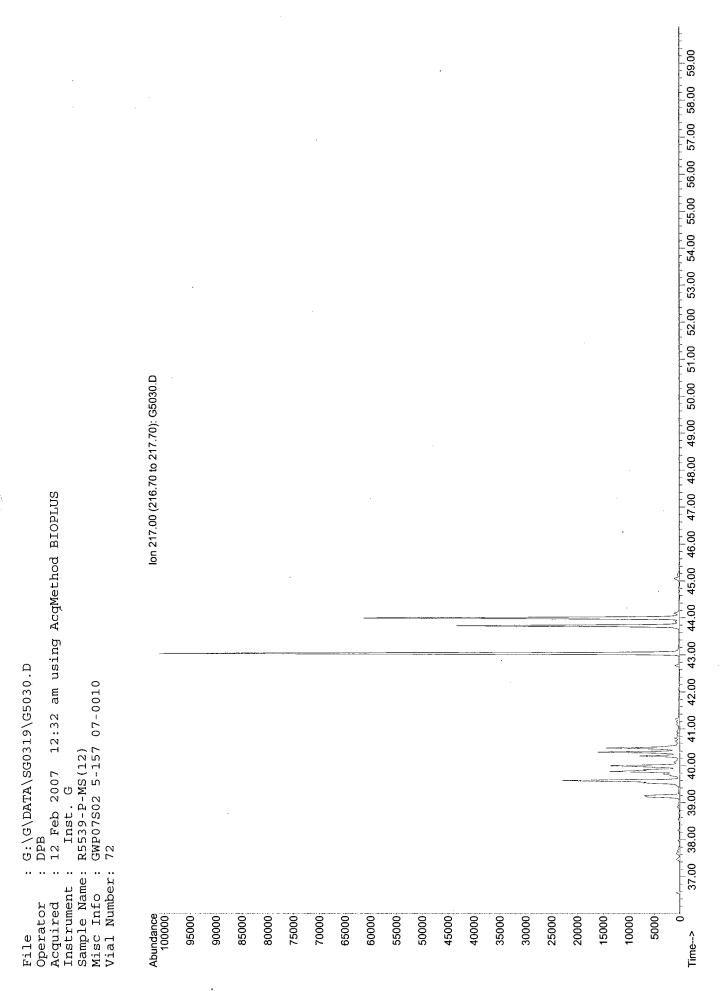




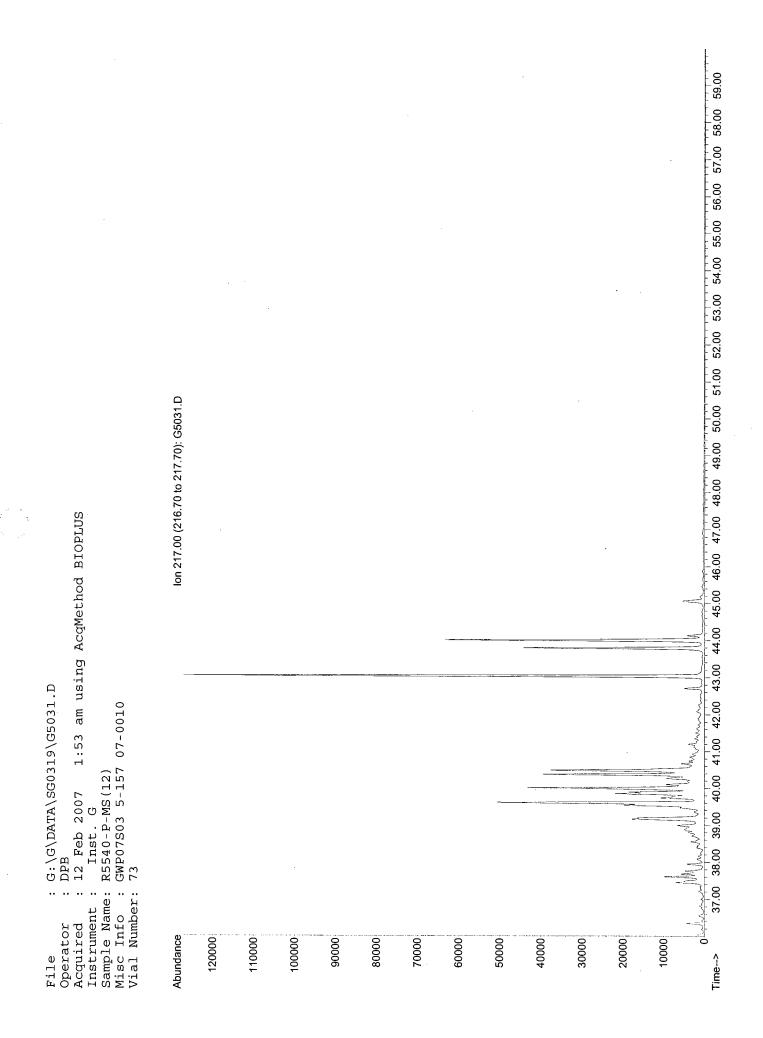


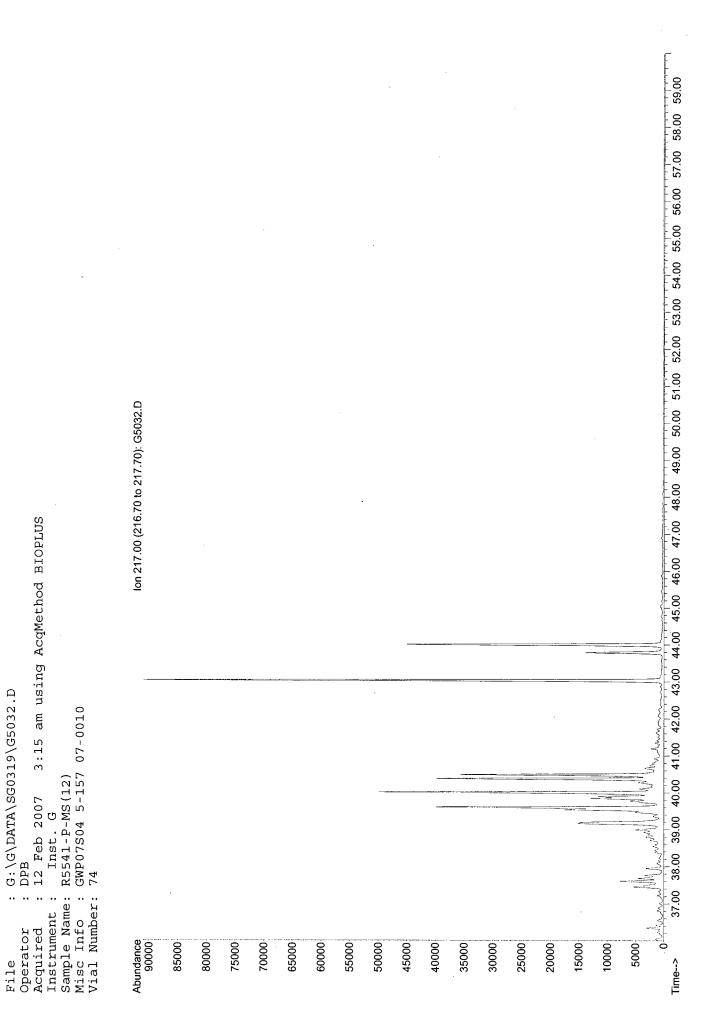


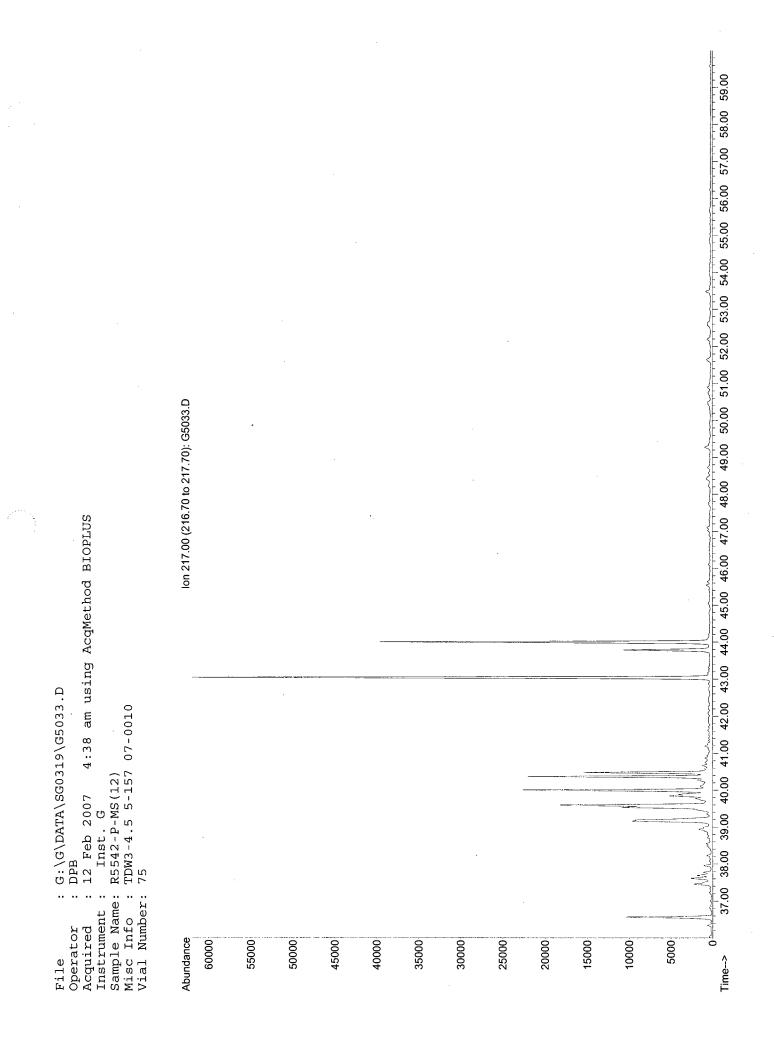


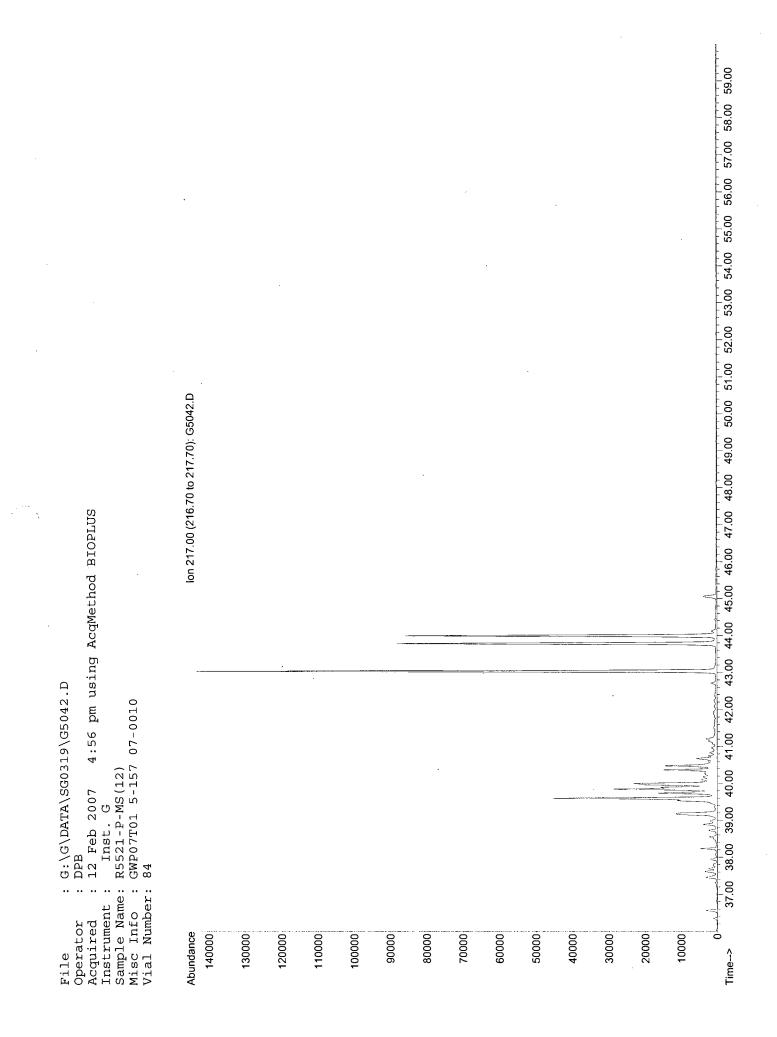


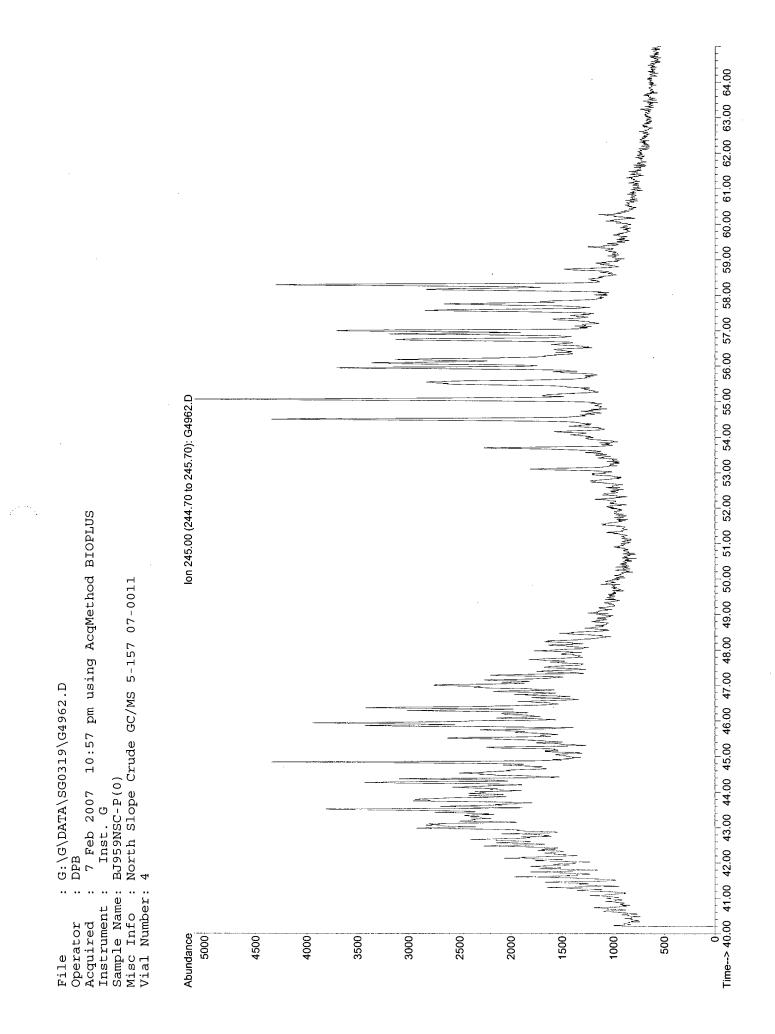
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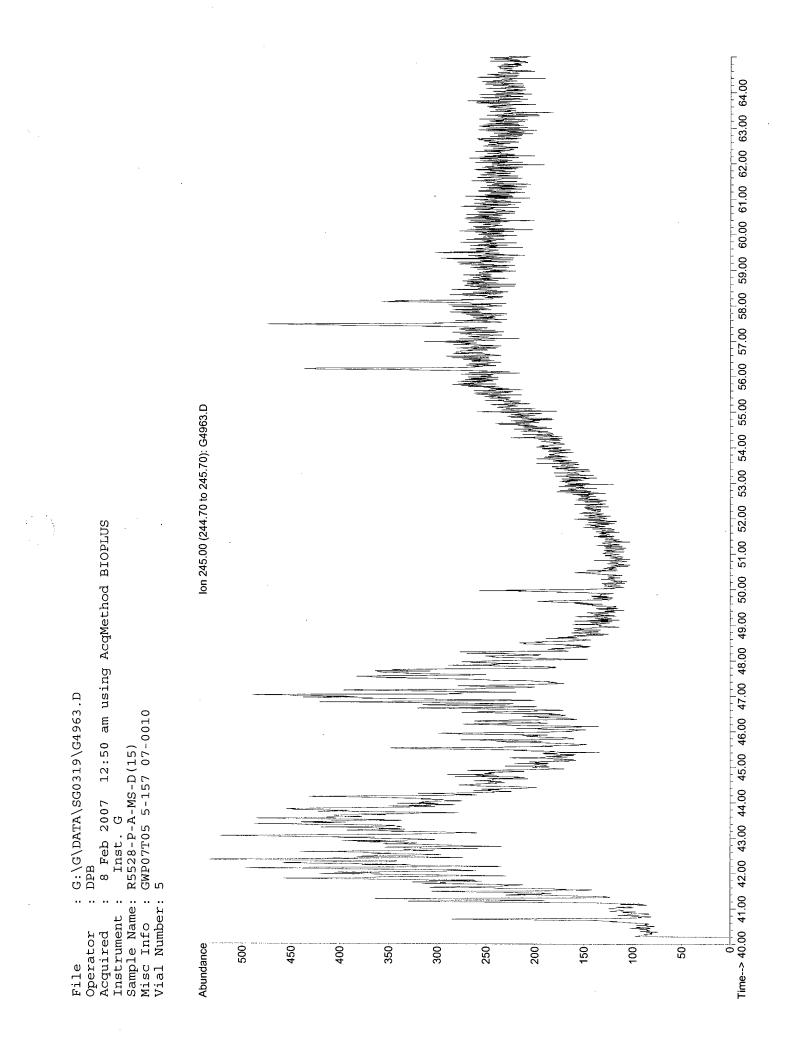


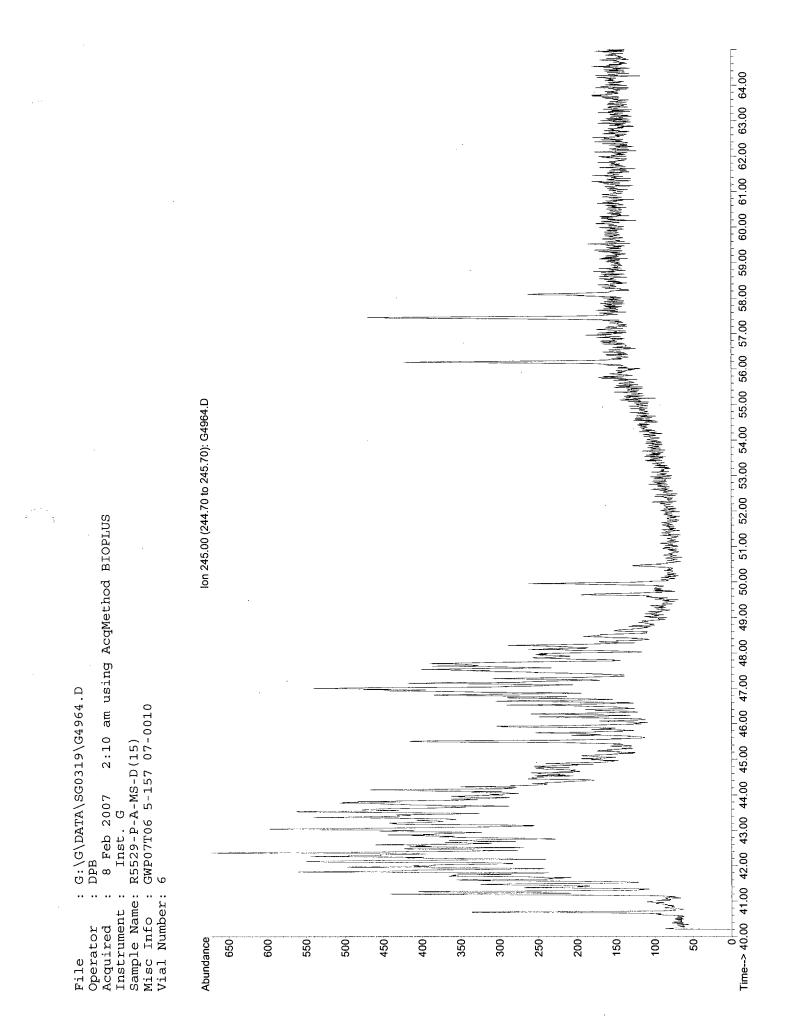




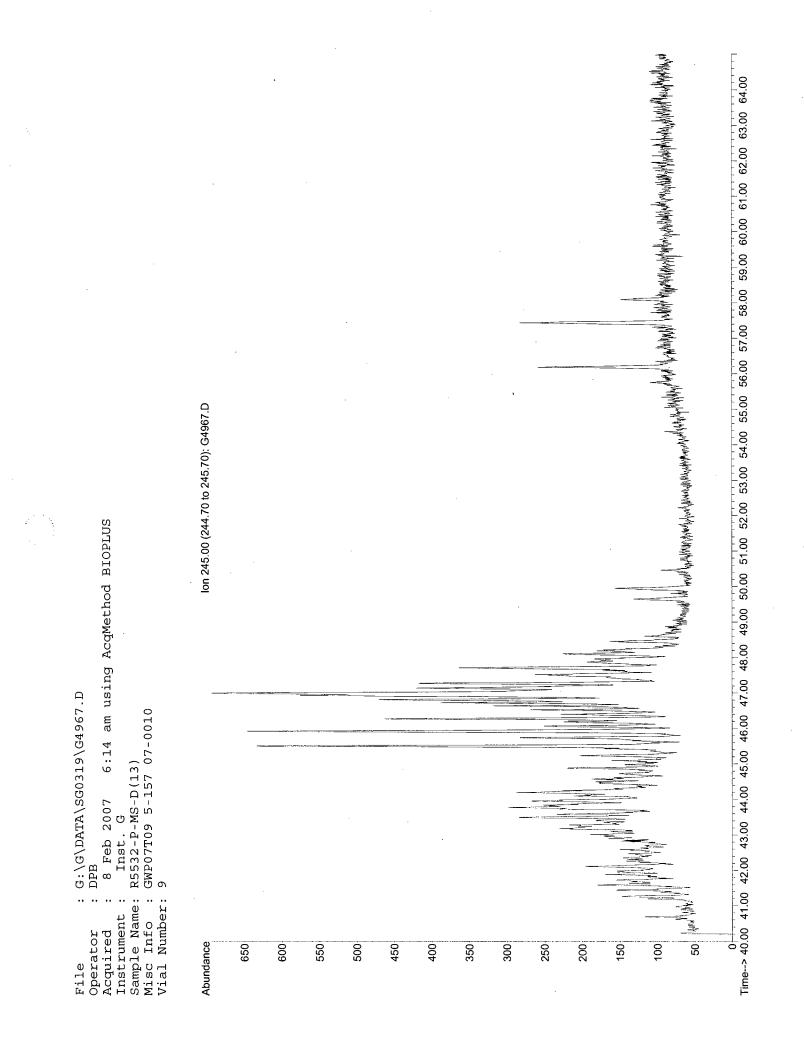


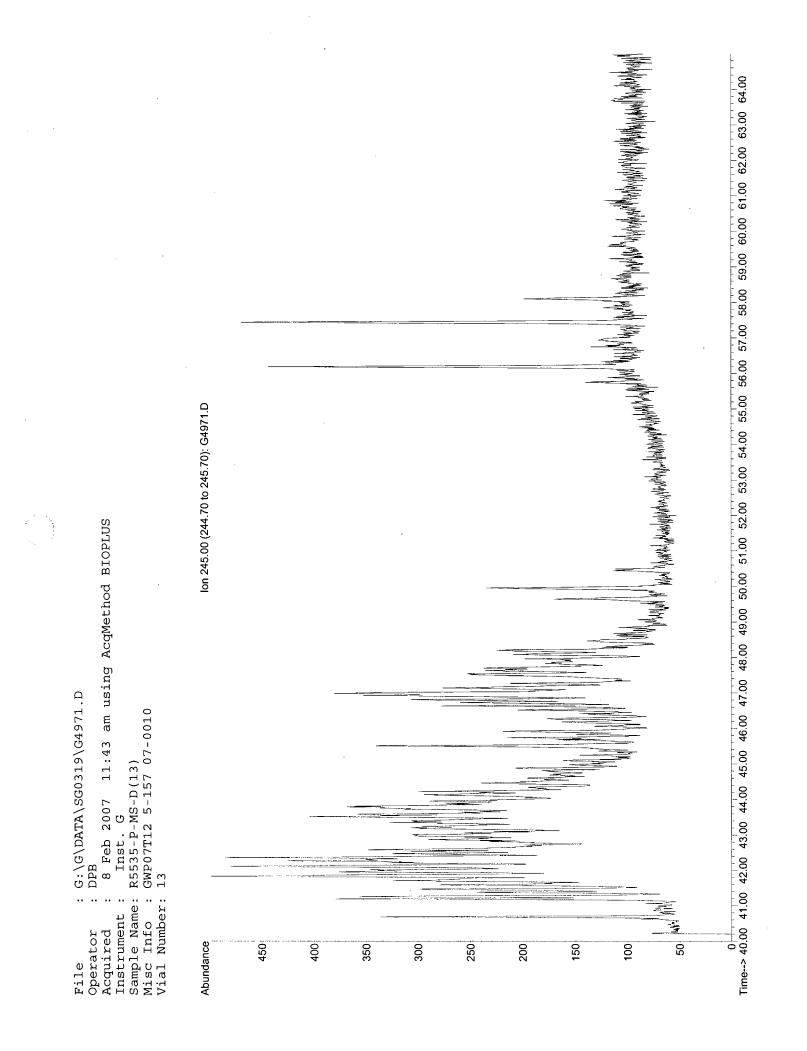


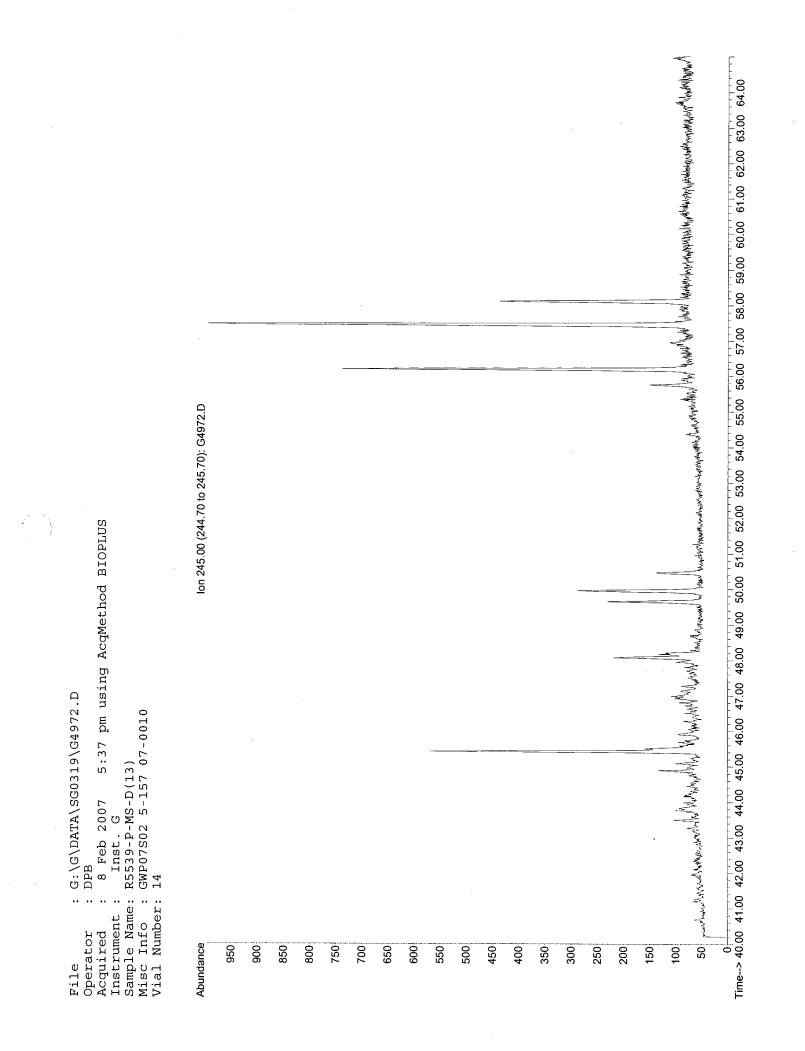


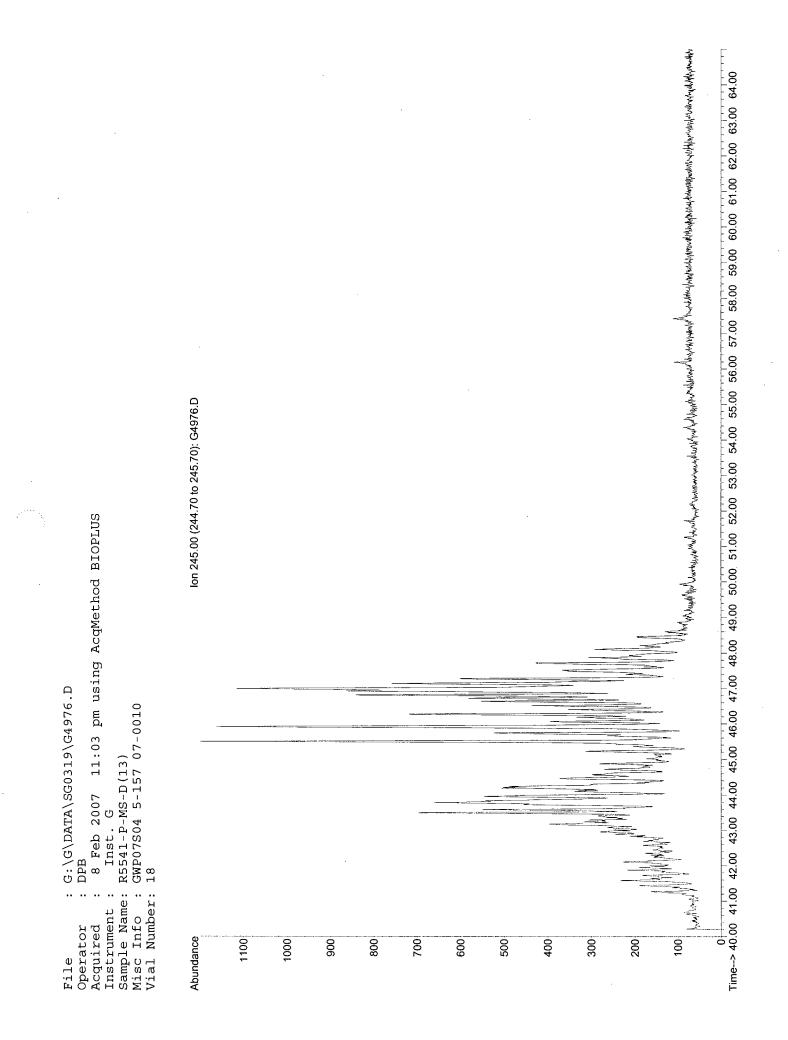


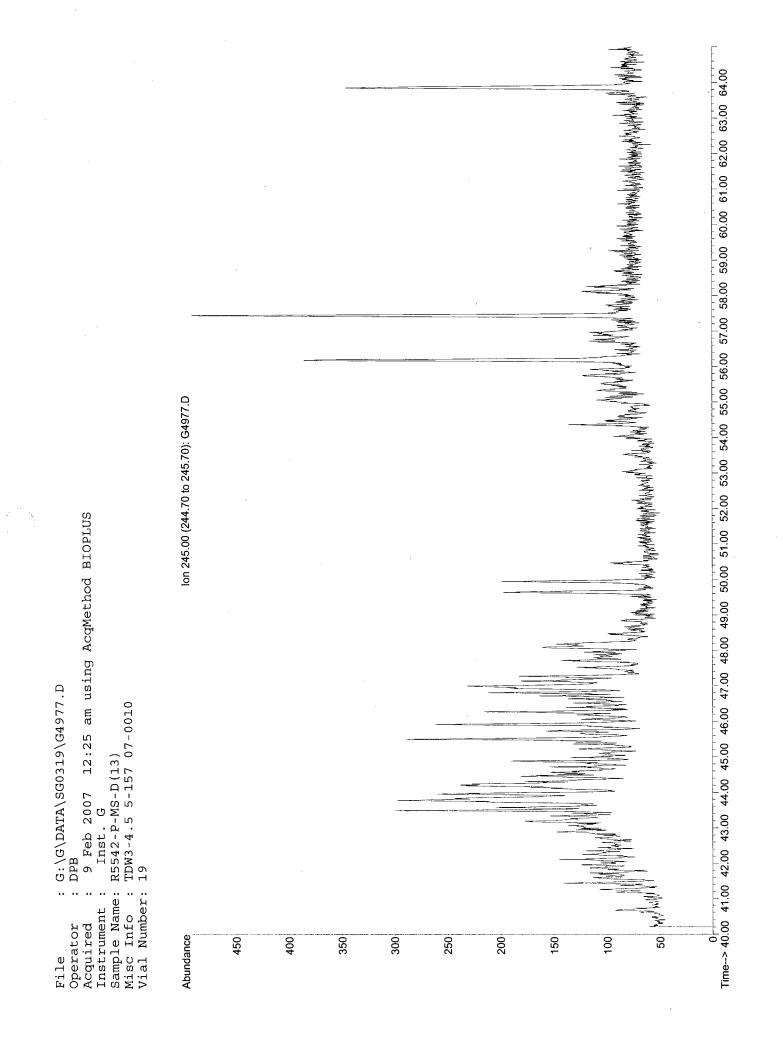


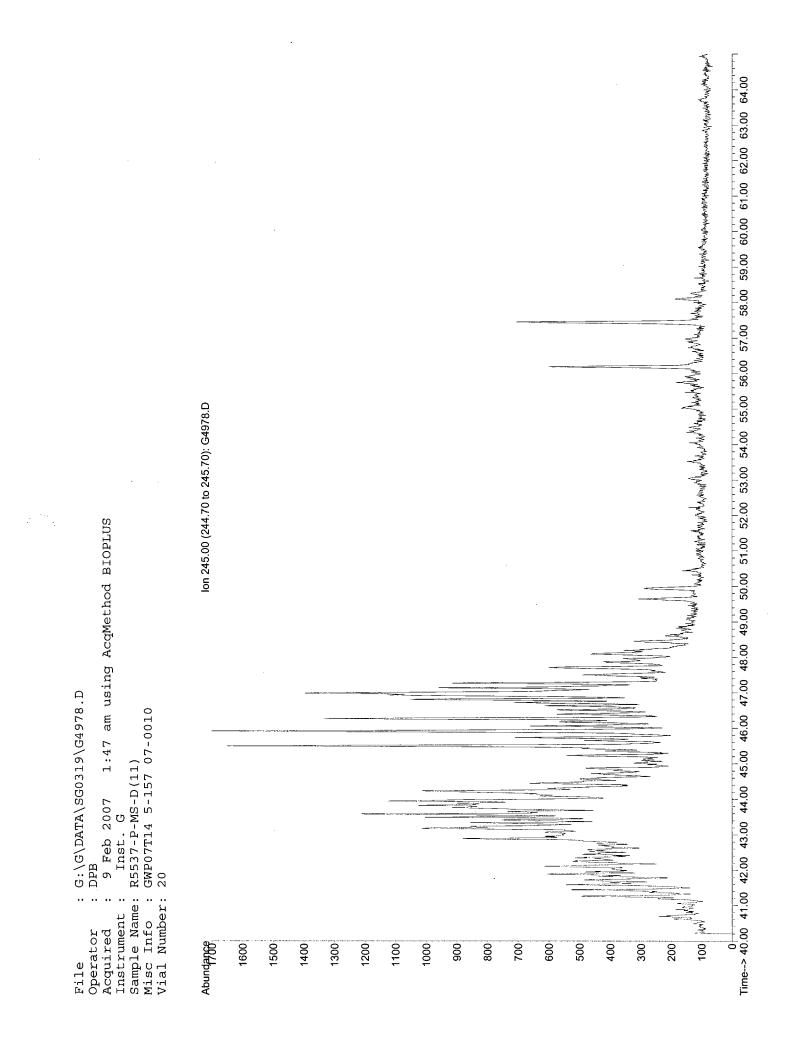


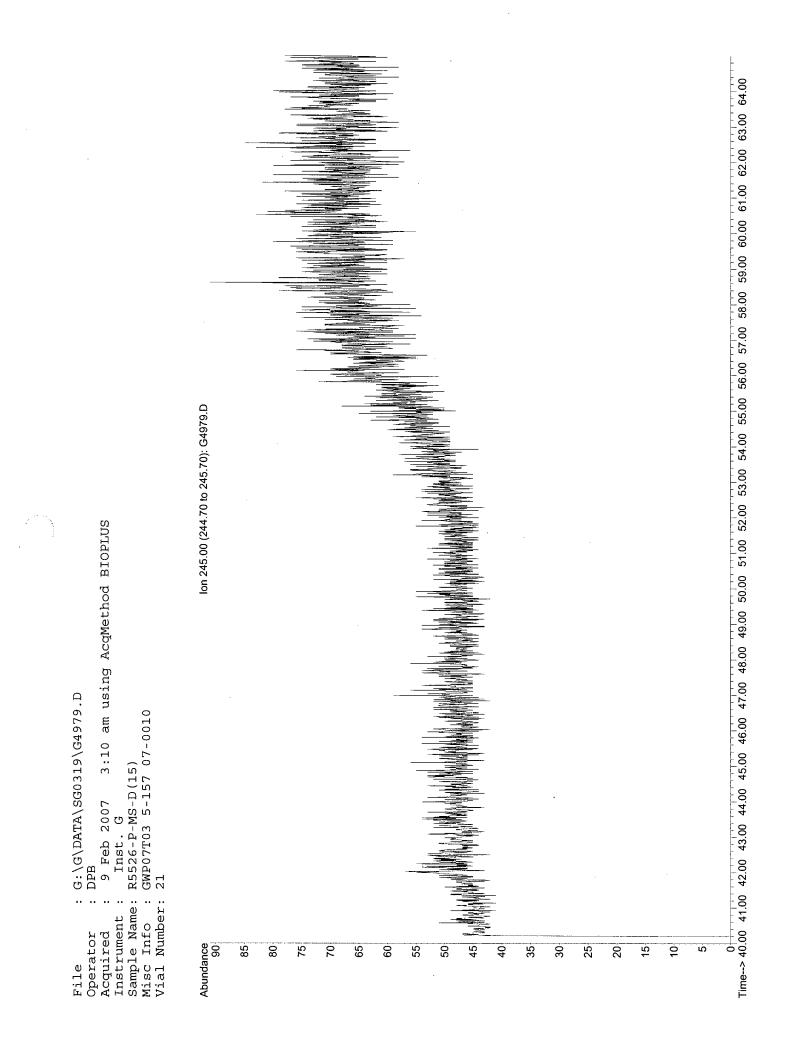


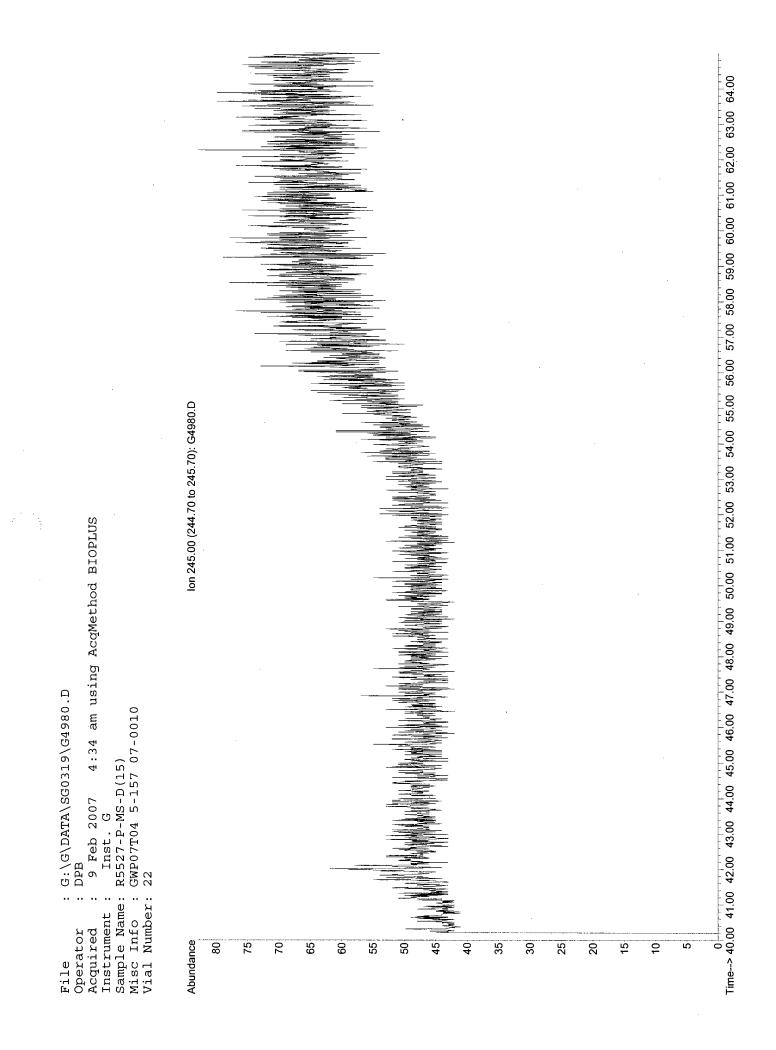


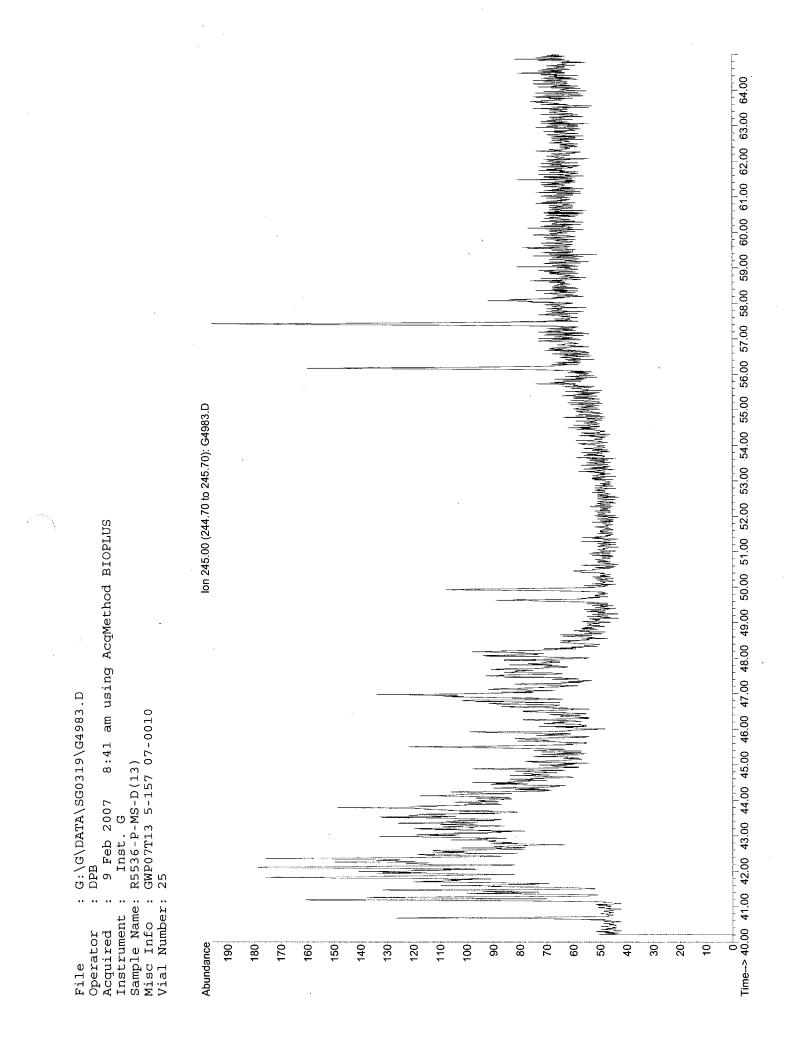


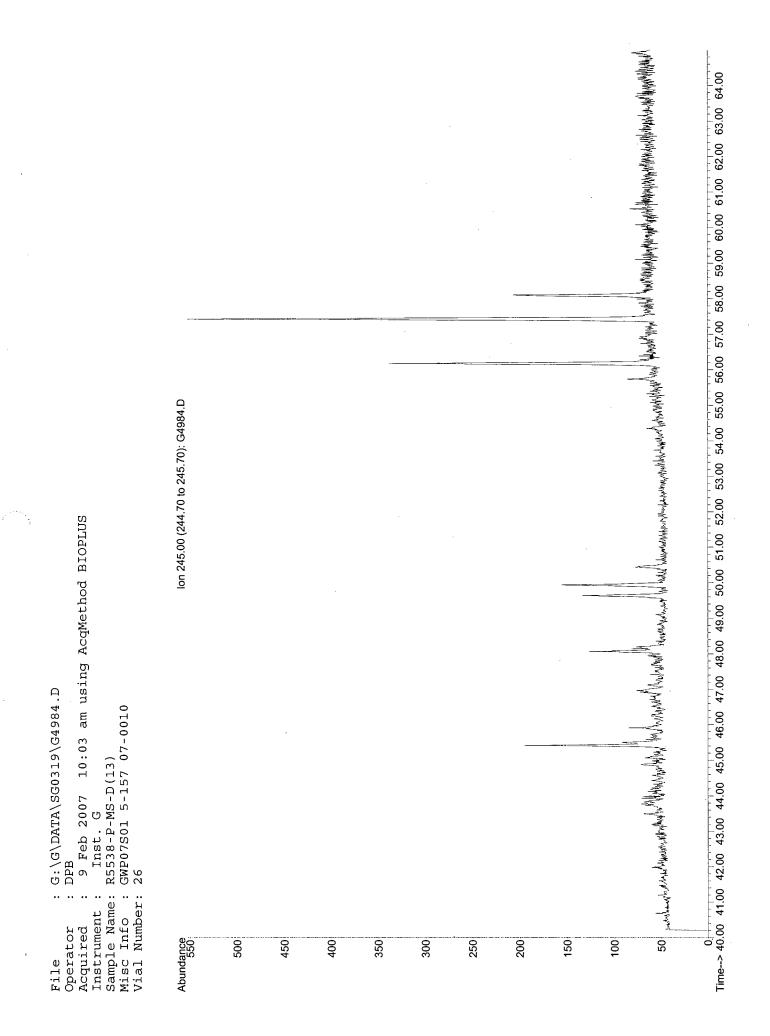


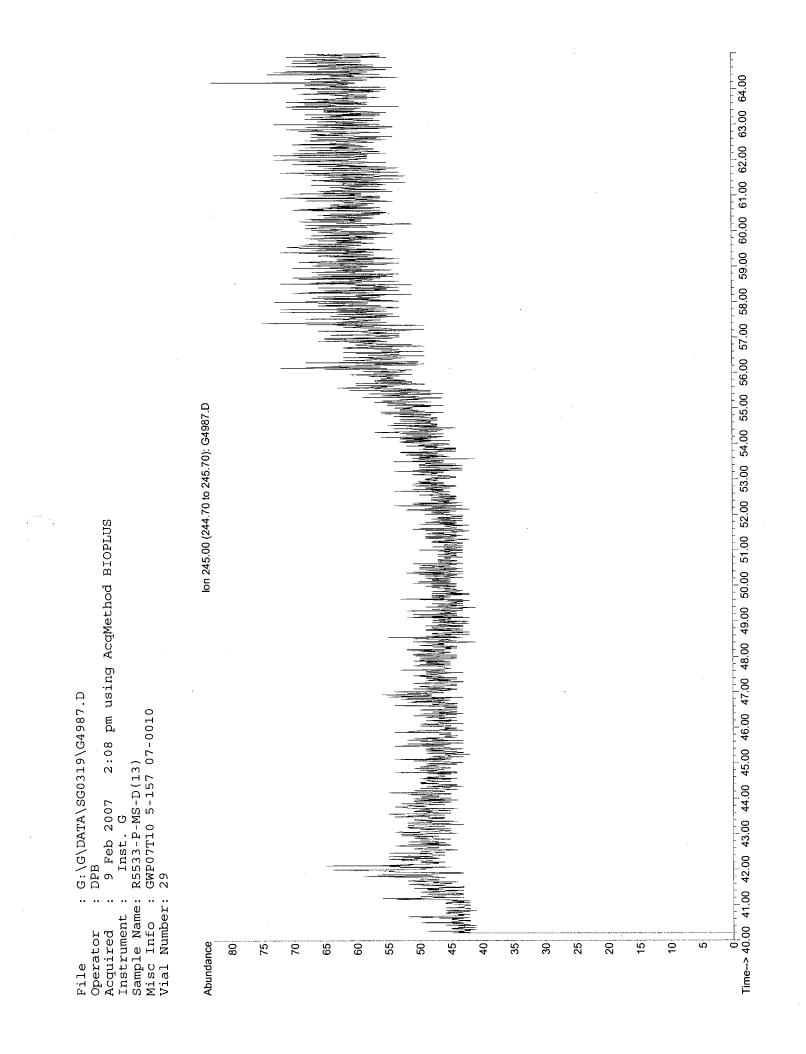


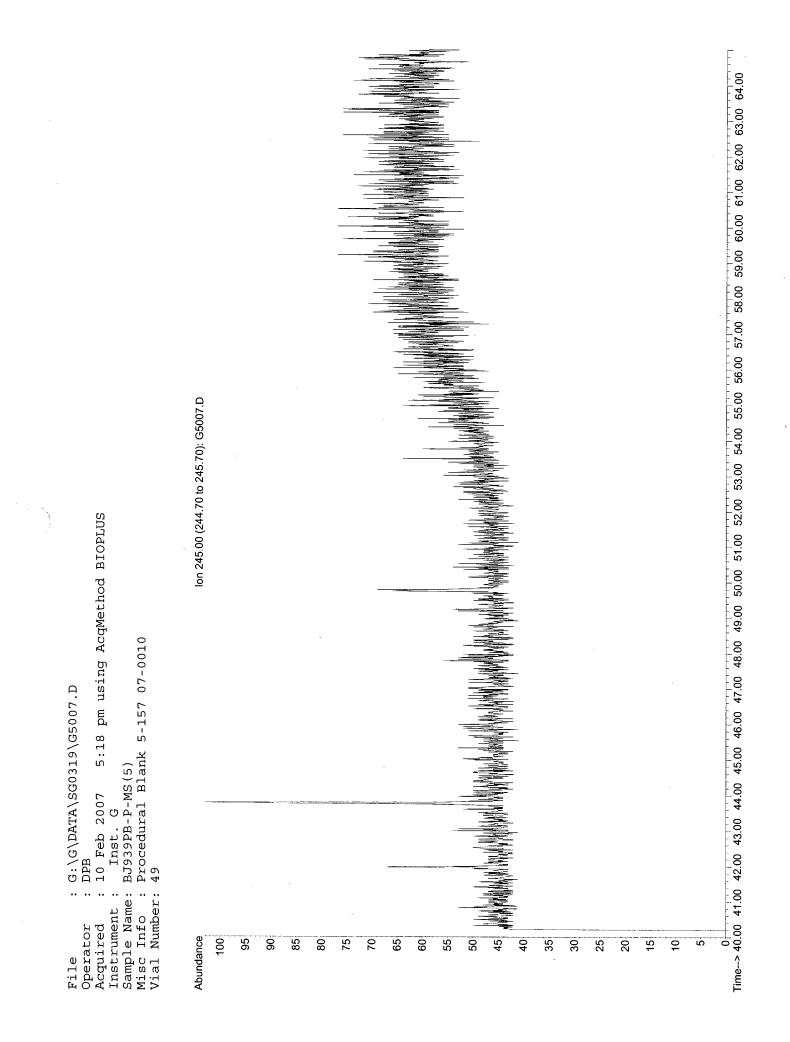


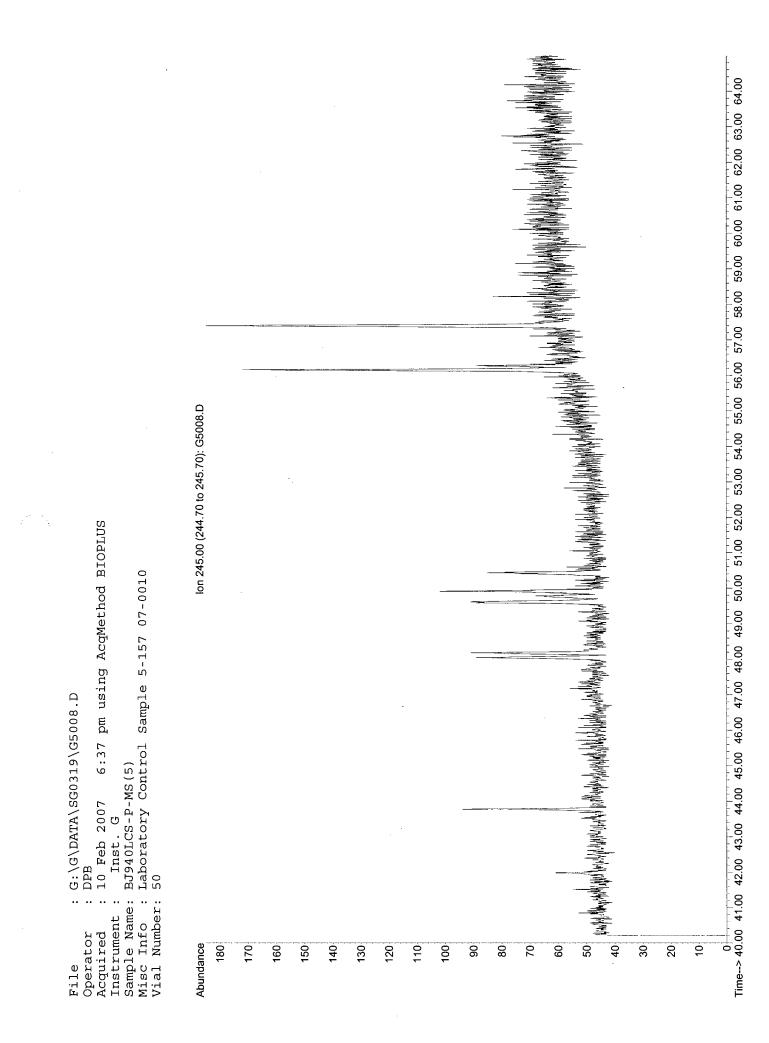


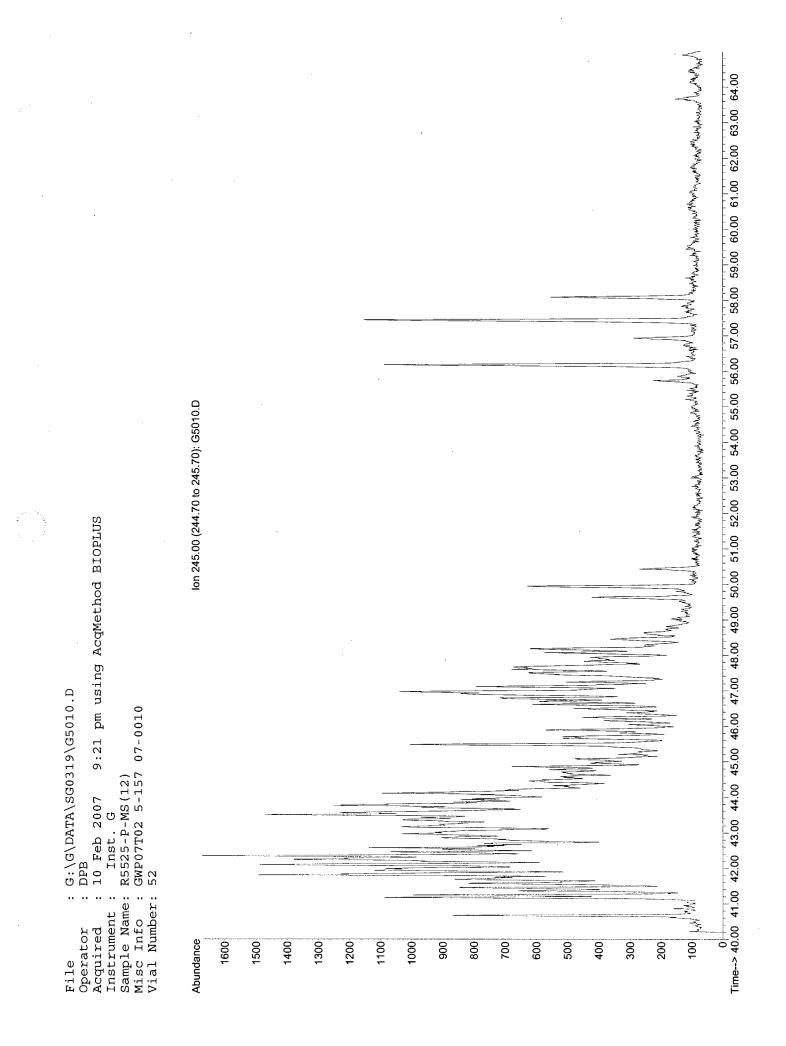


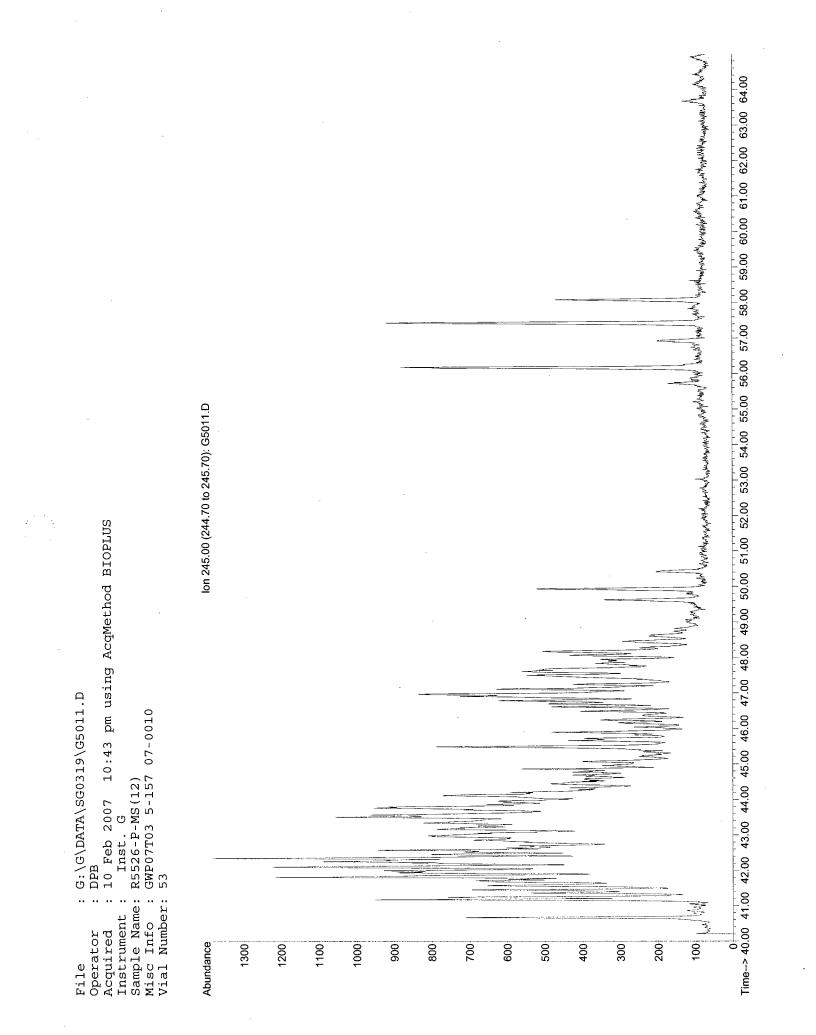


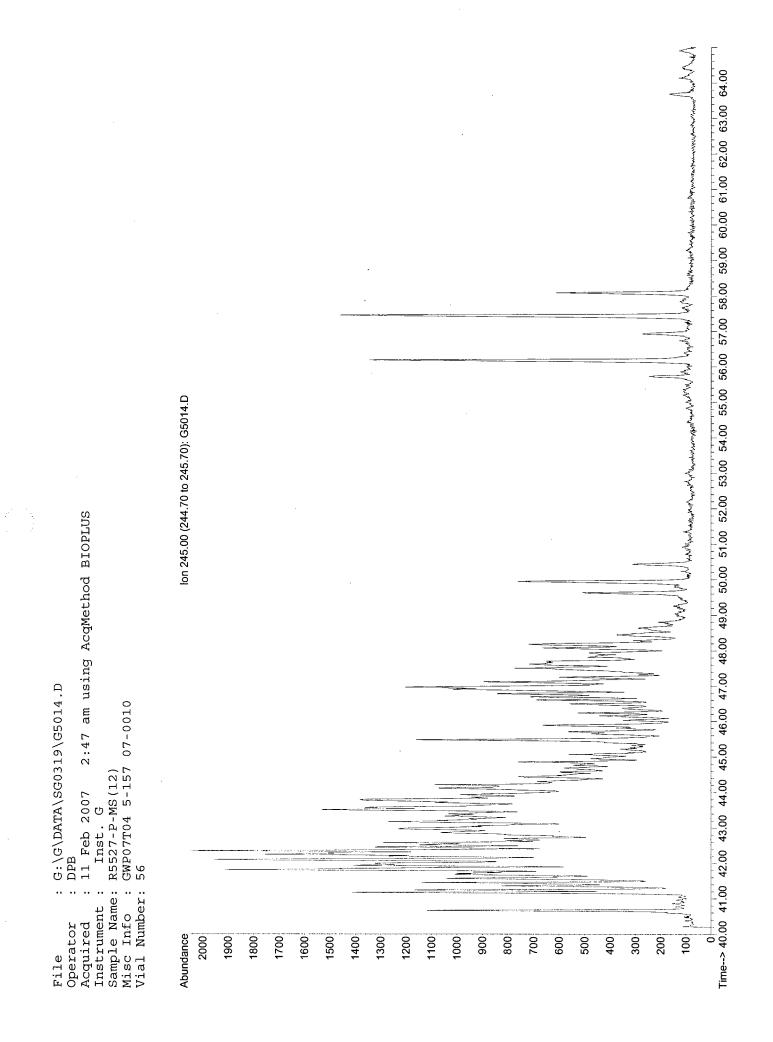


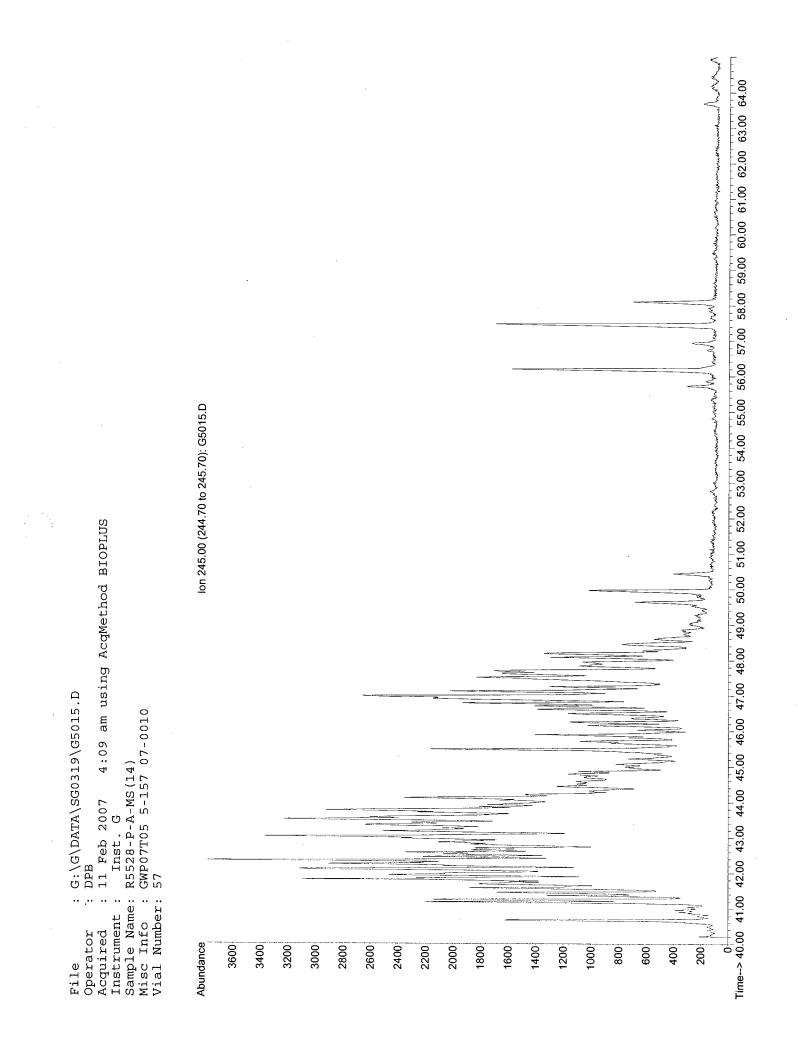


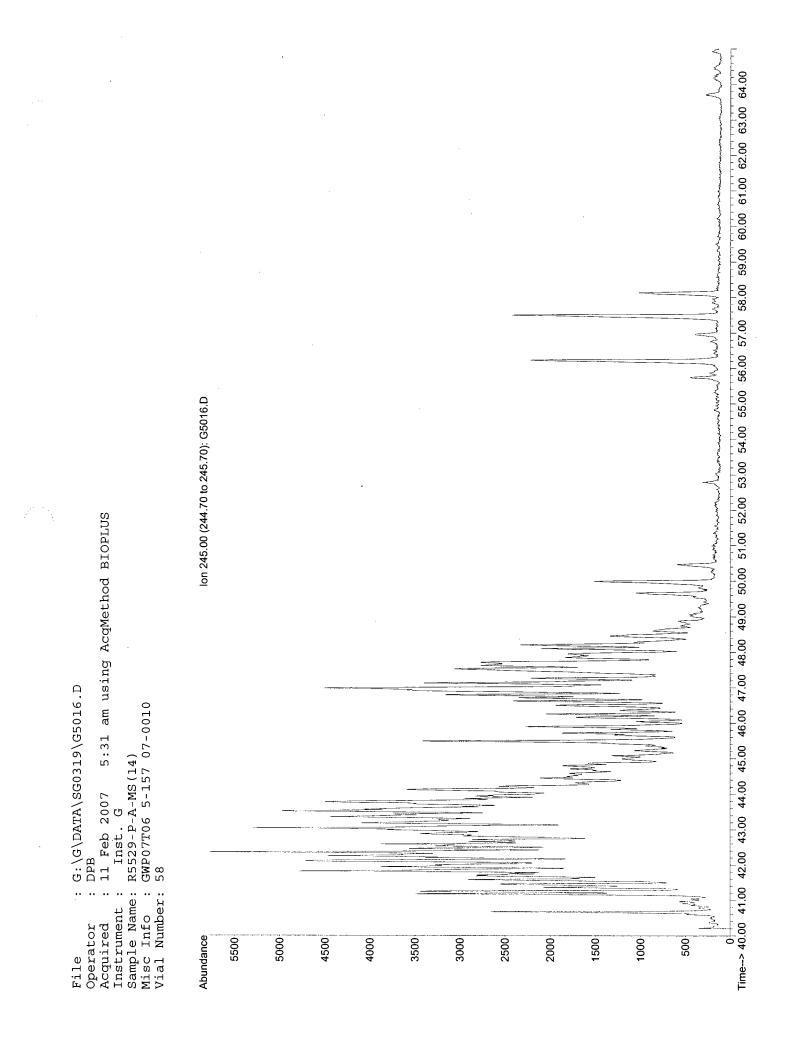


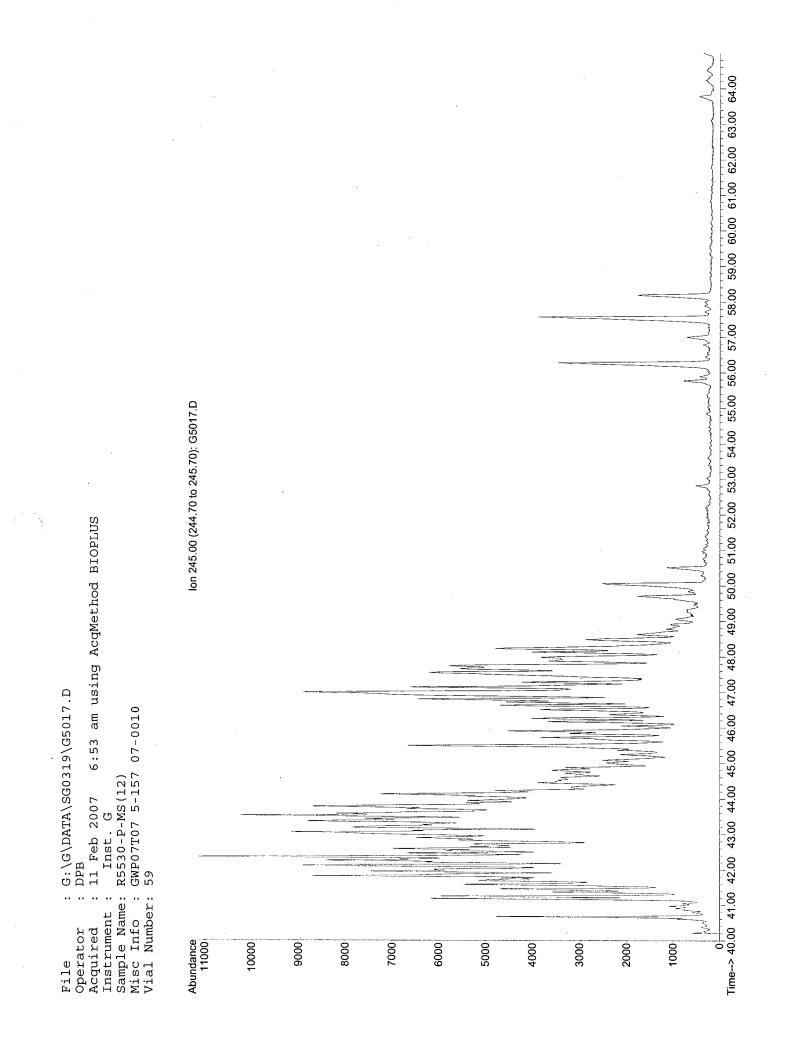


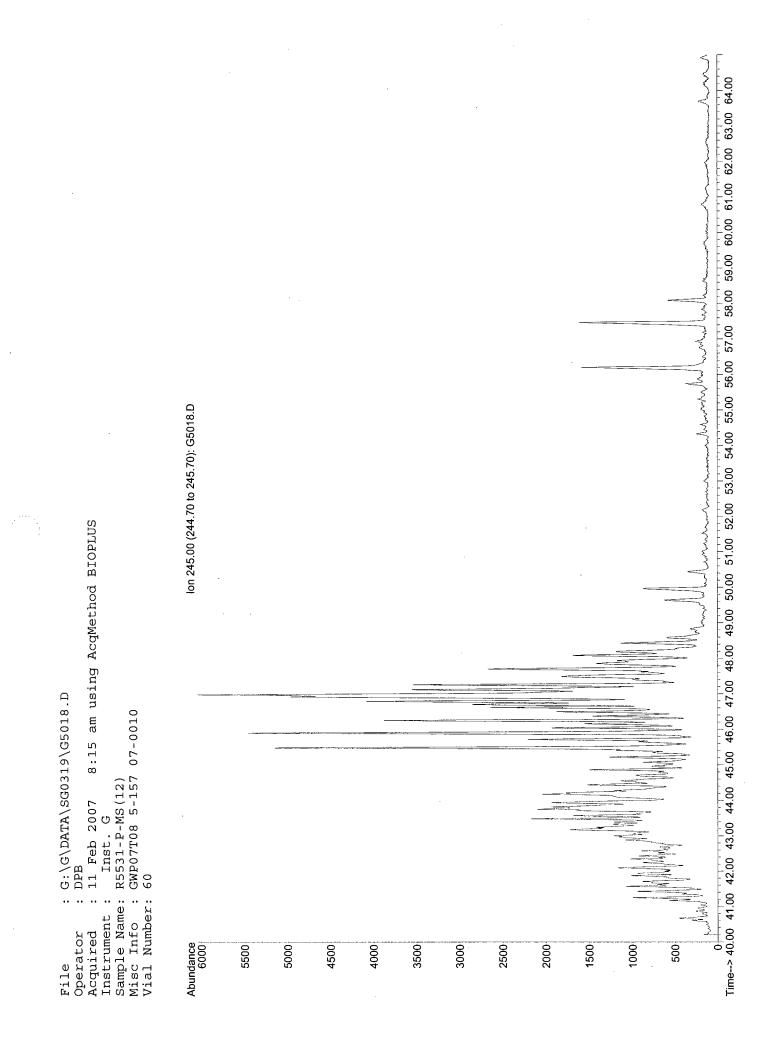


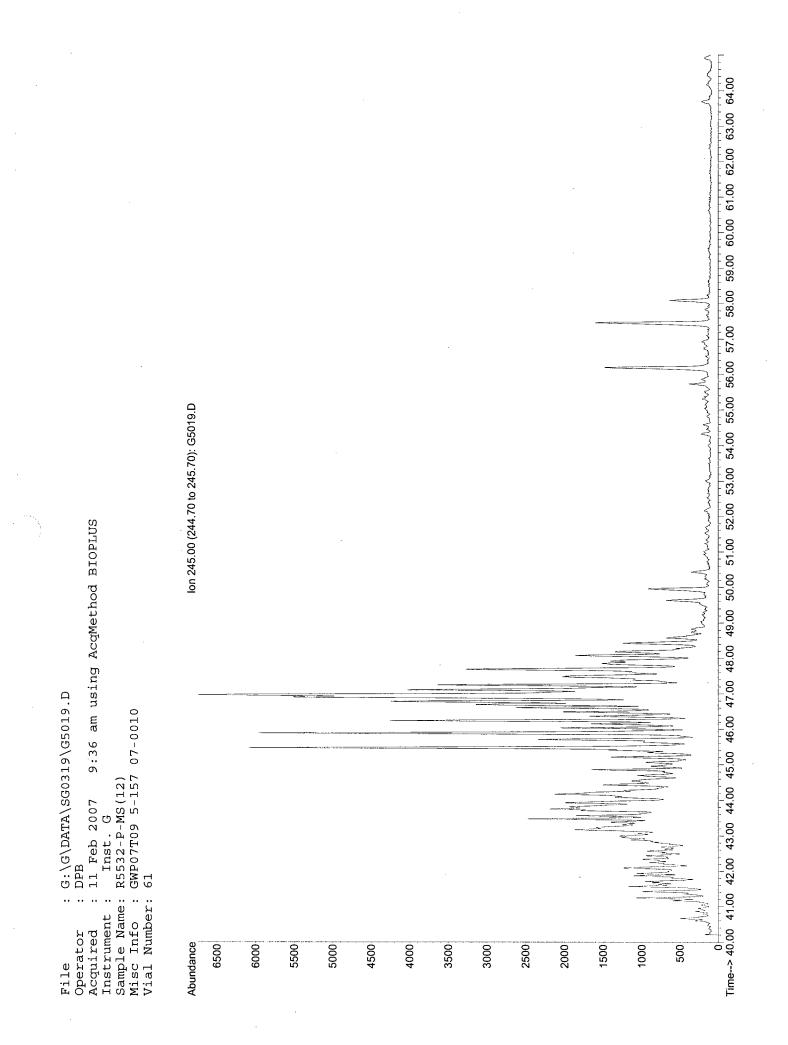


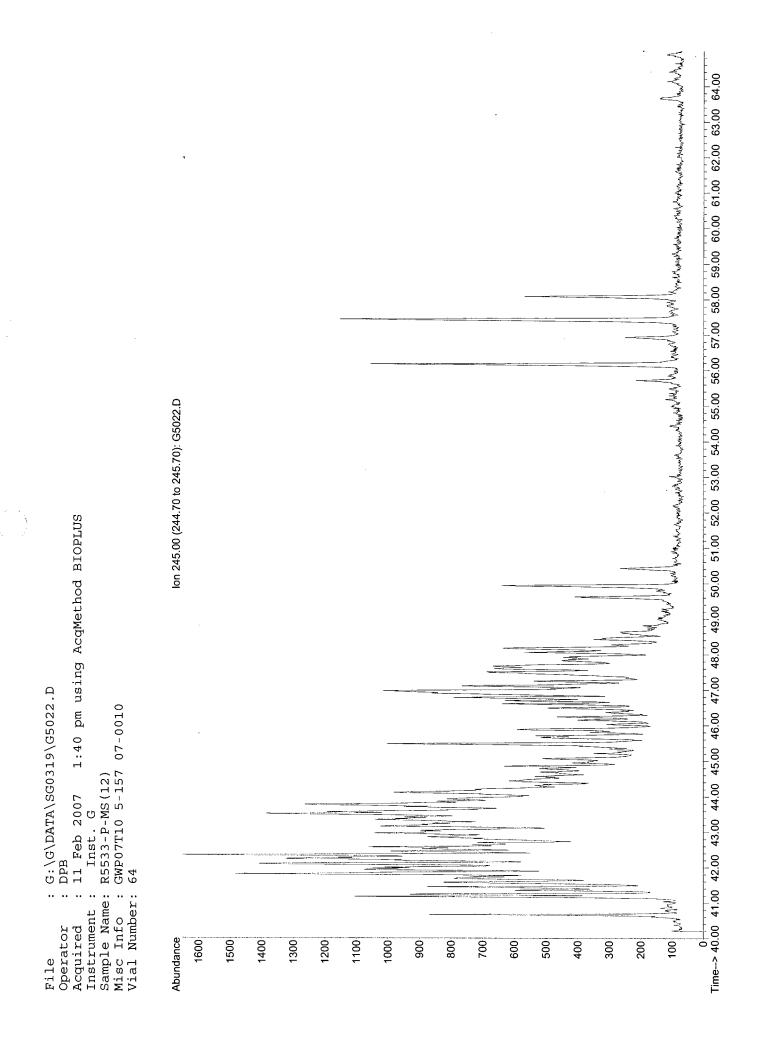




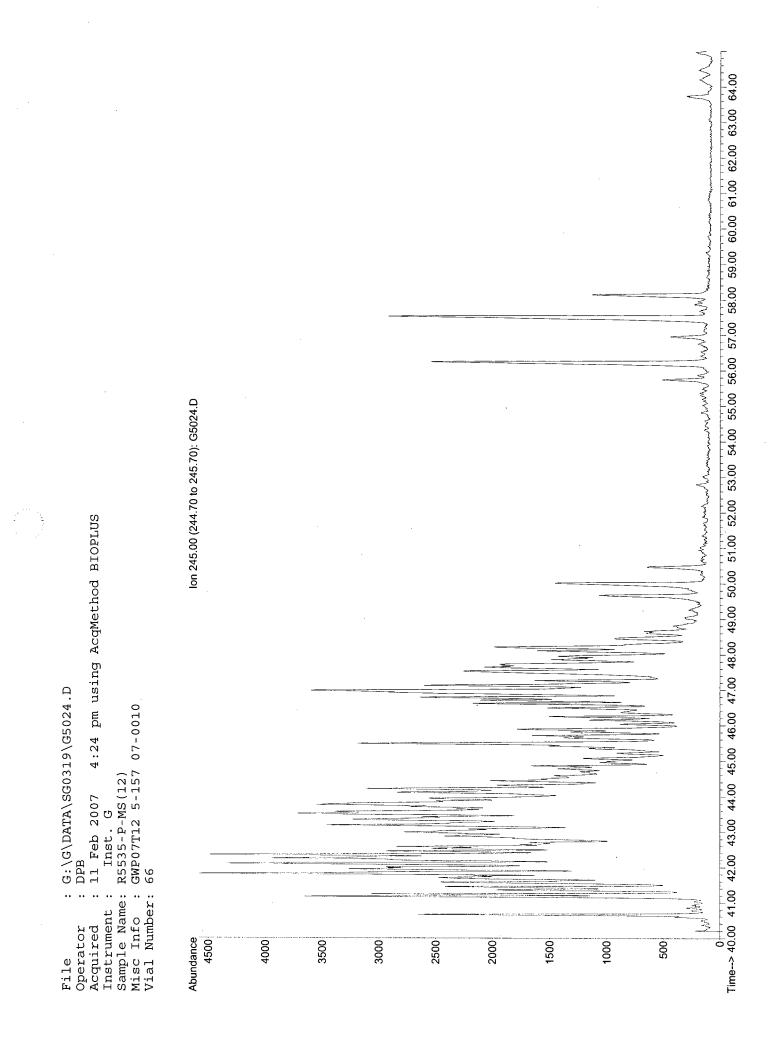


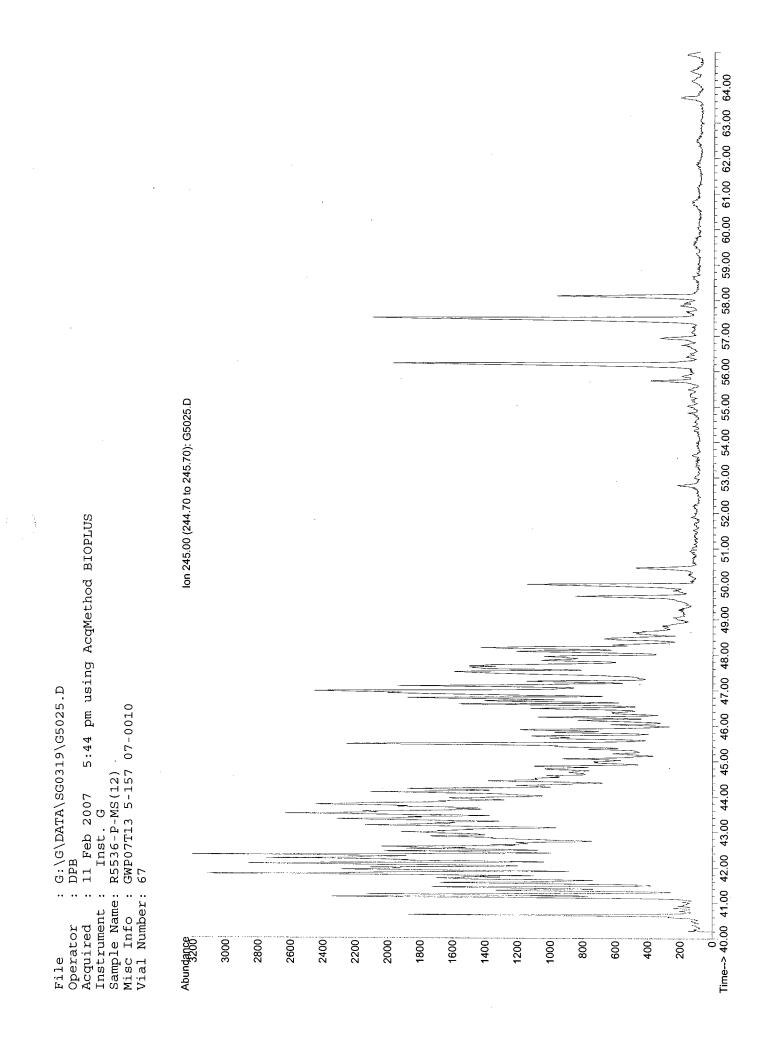


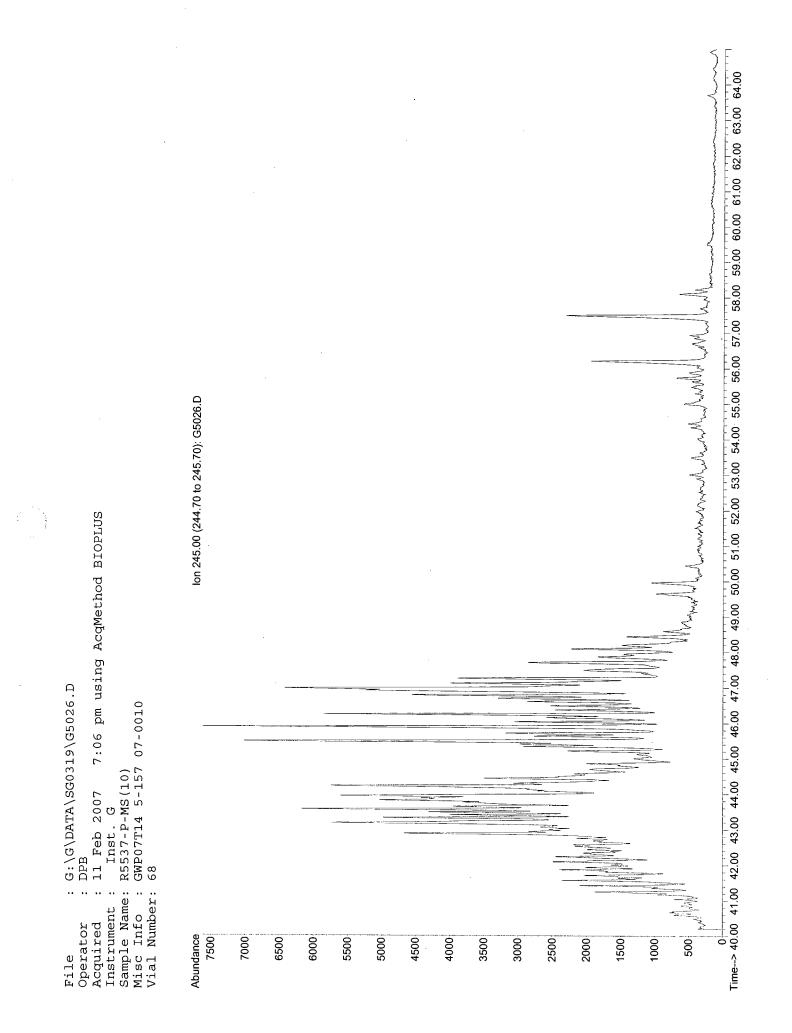


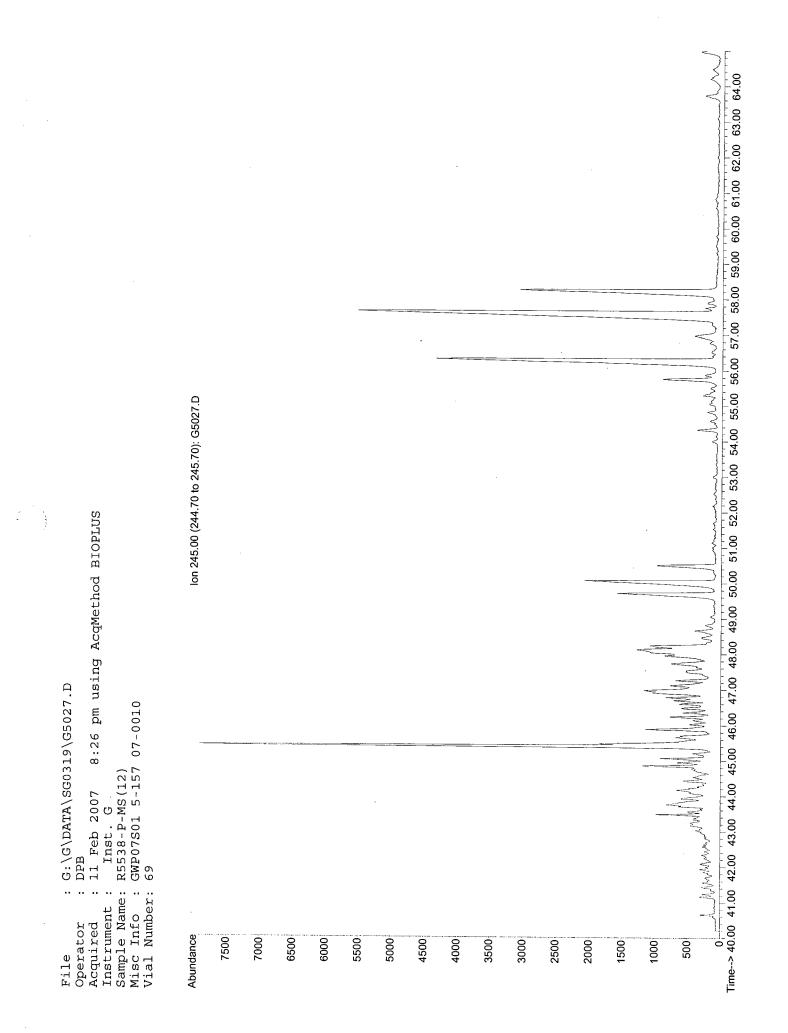


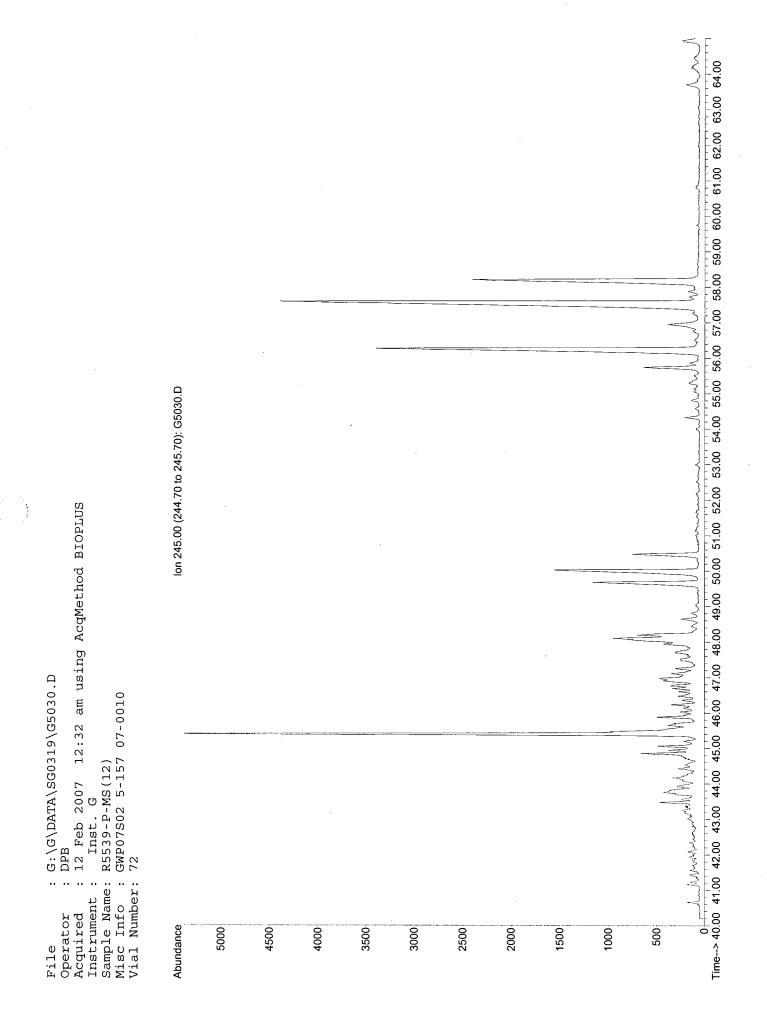


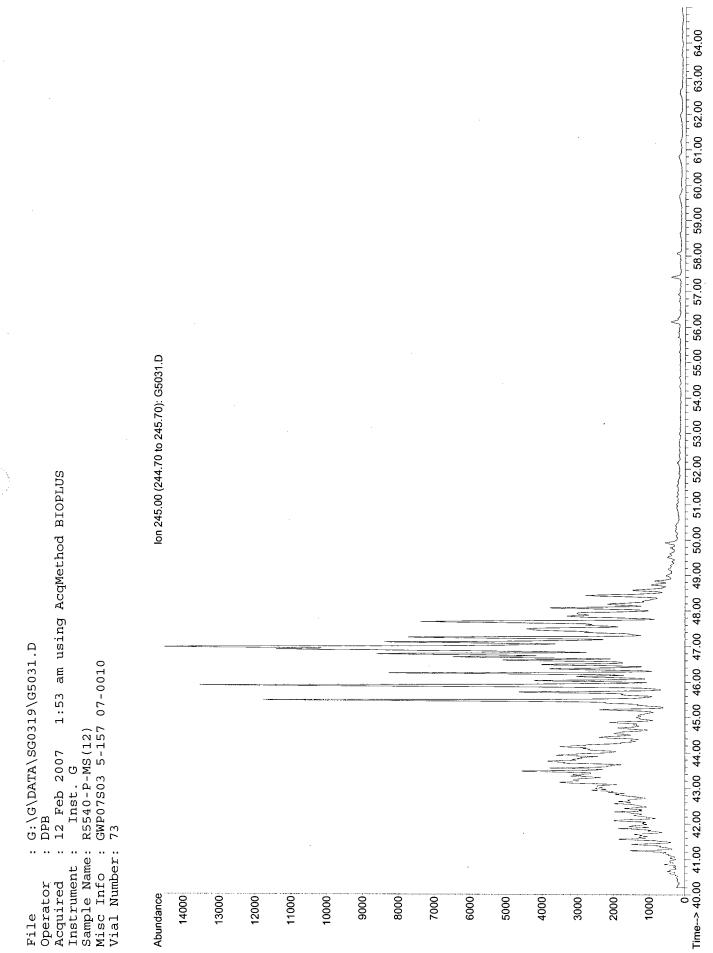




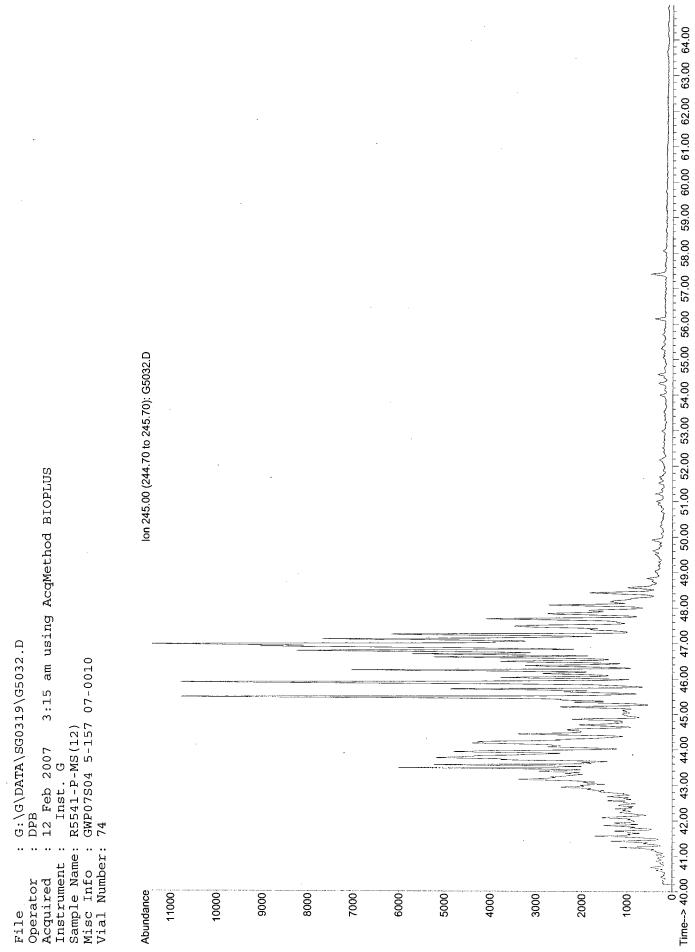






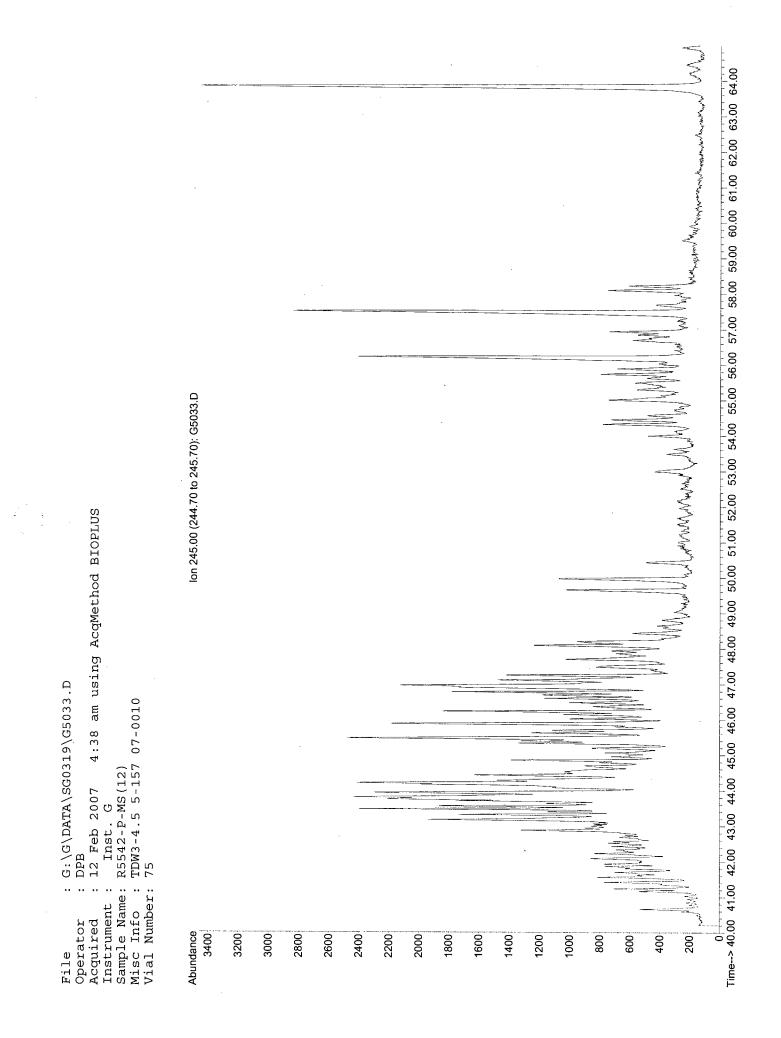


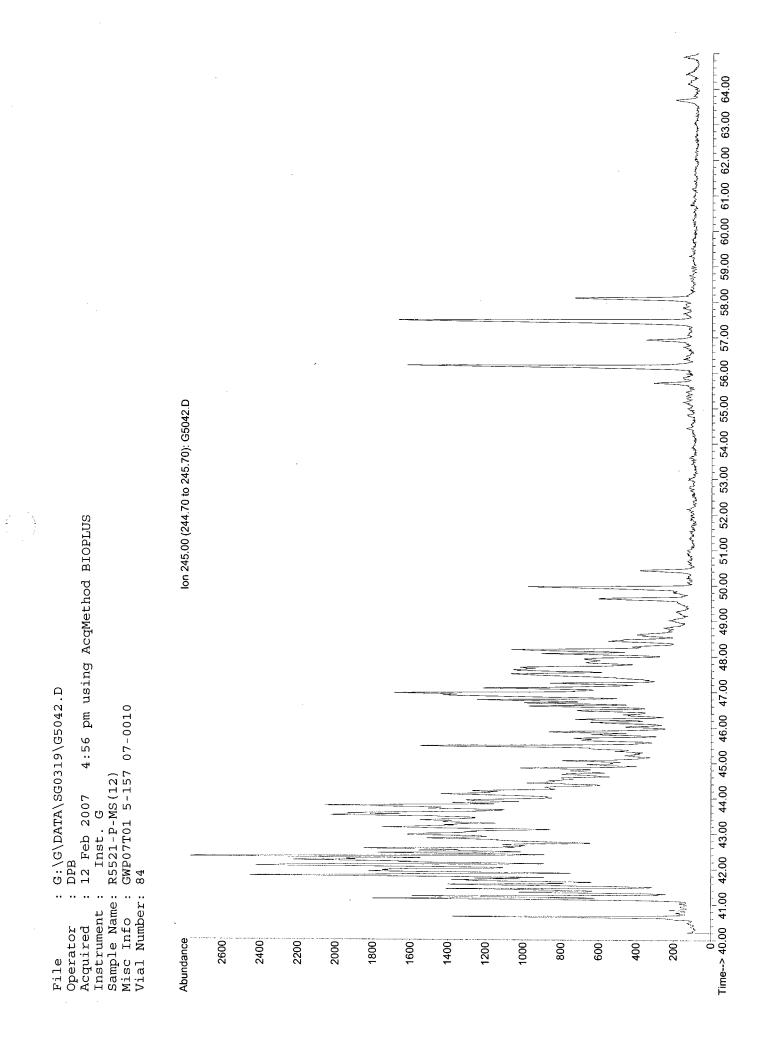
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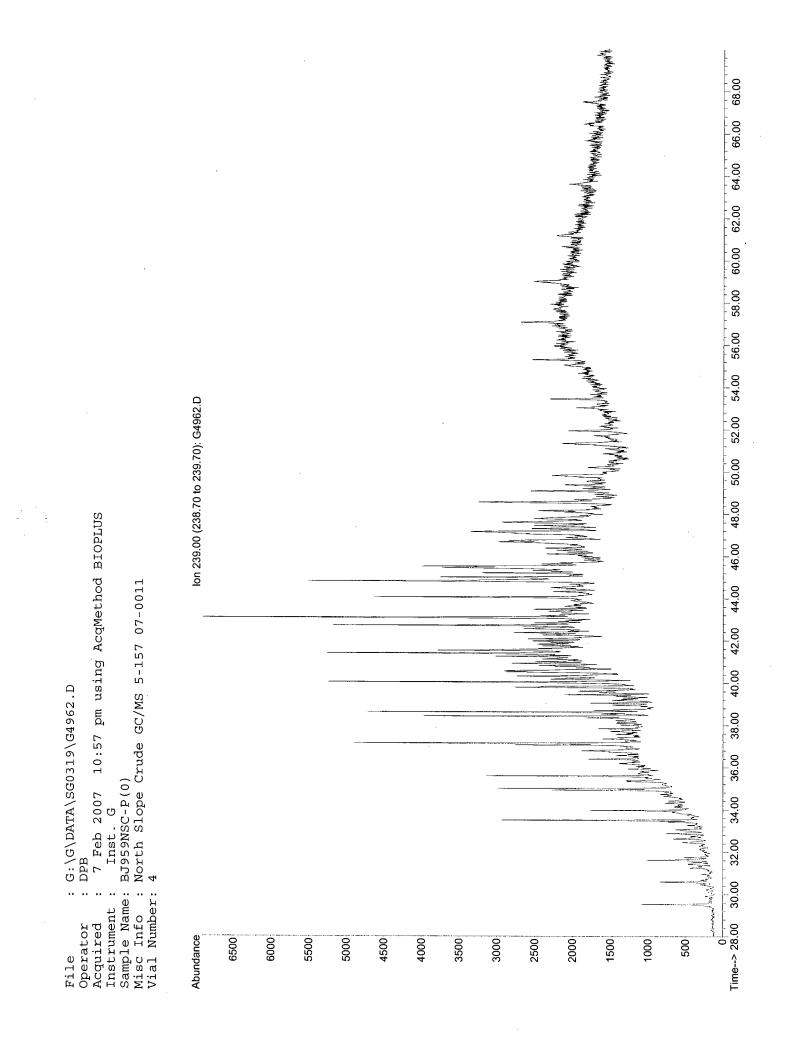


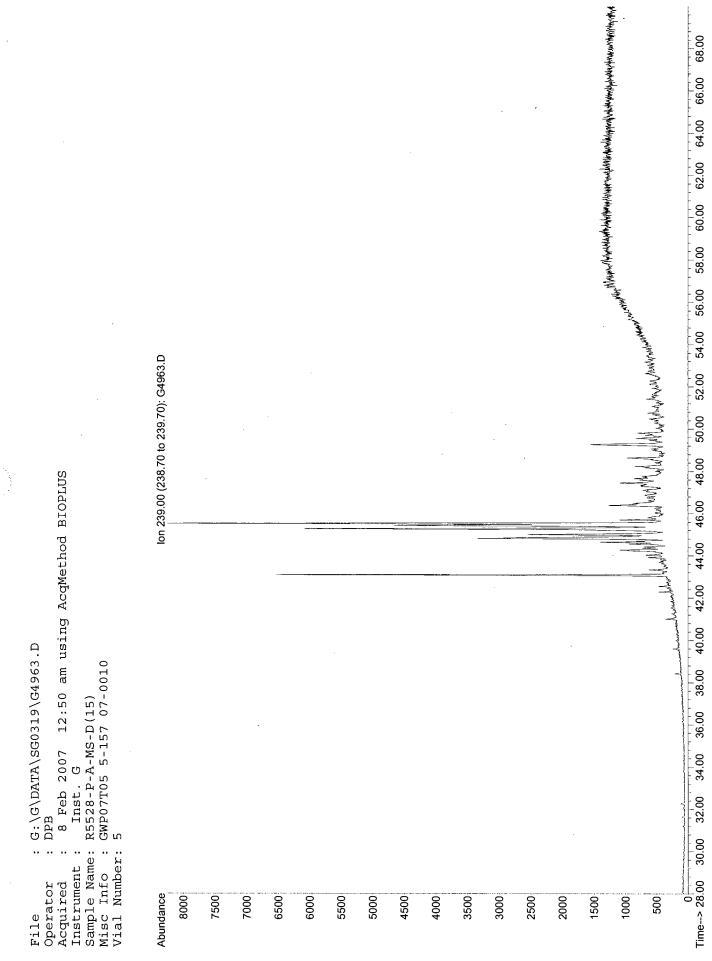
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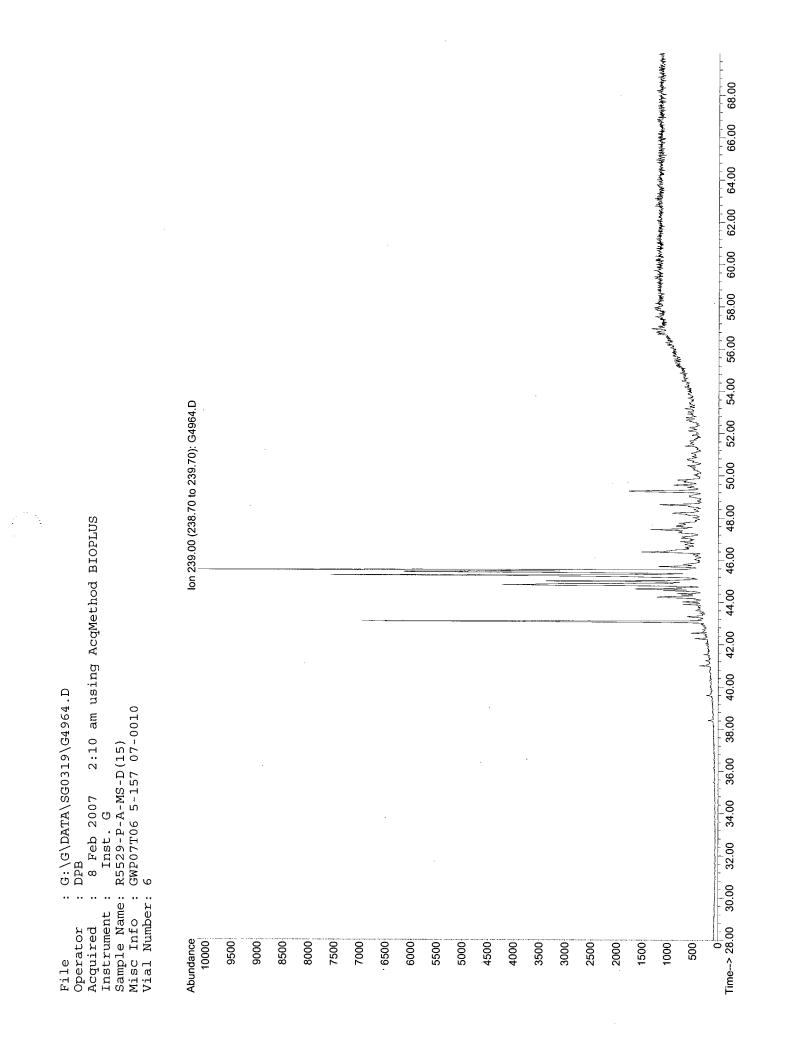
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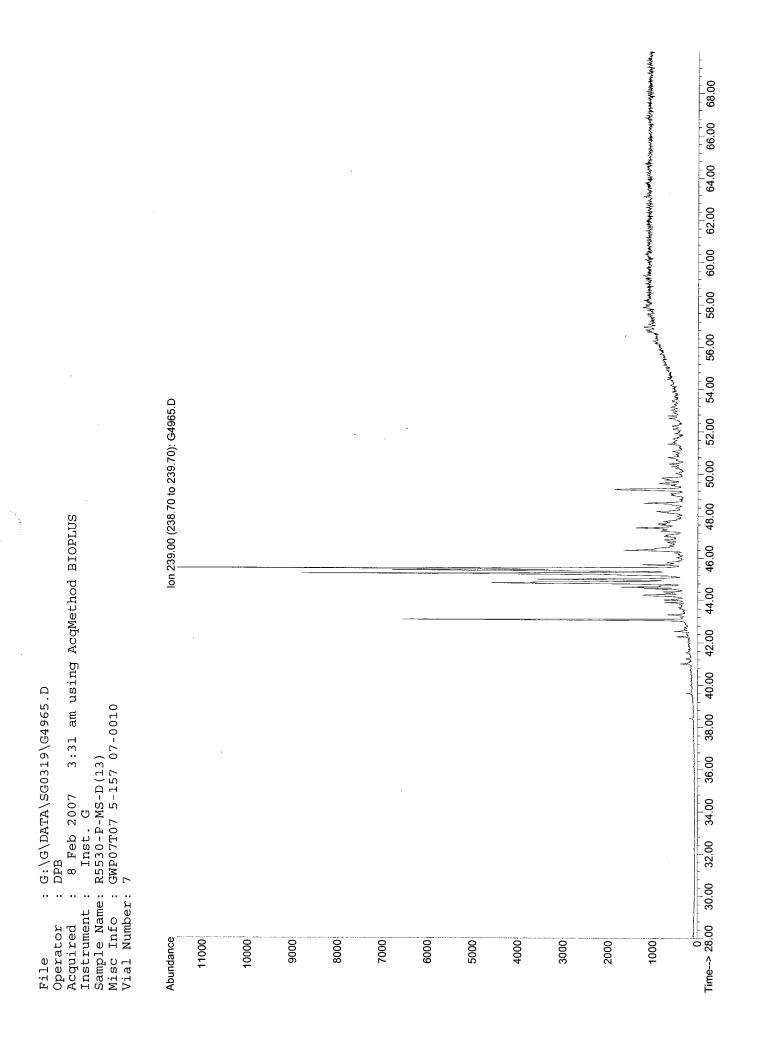


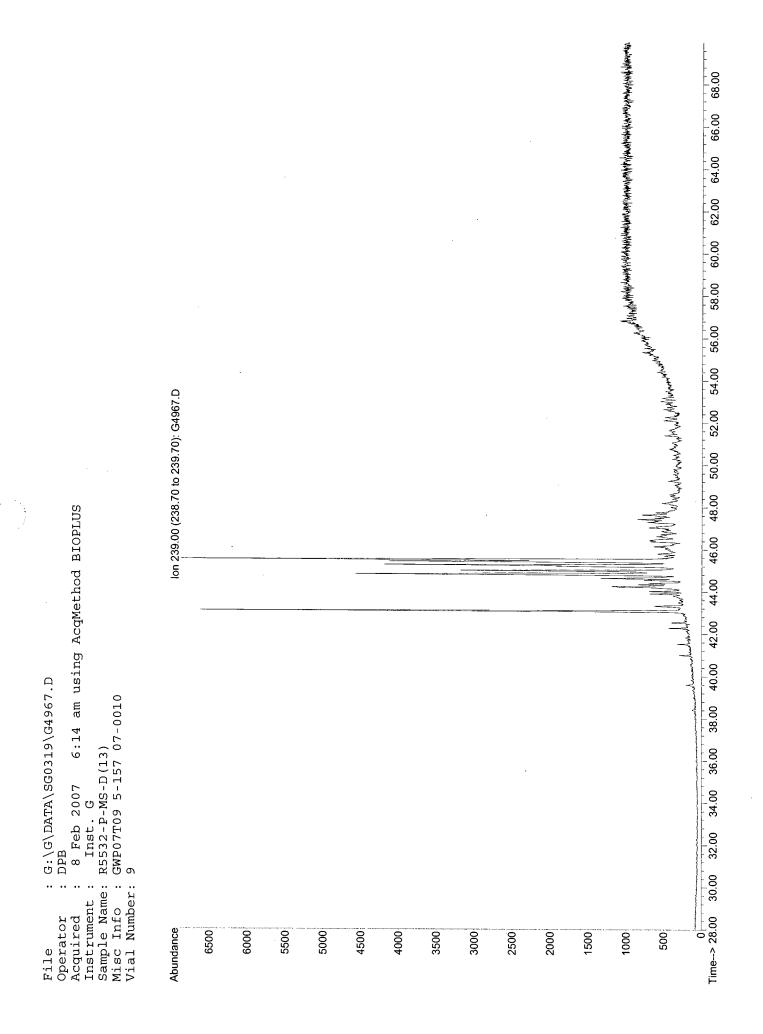


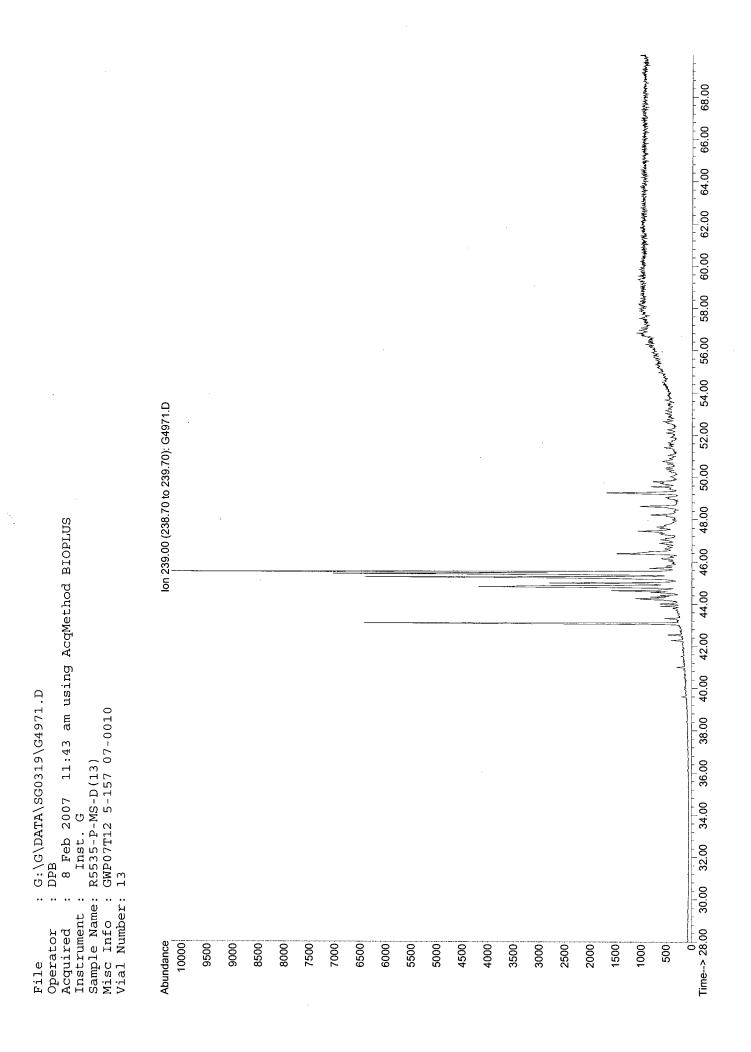












Ion 239.00 (238.70 to 239.70): G4972.D 5:37 pm using AcqMethod BIOPLUS : G:\G\DATA\SG0319\G4972.D : DPB Acquired : 8 Feb 2007 5:37 pm 1 Instrument : Inst. G Sample Name: R5539-P-MS-D(13) Misc Info : GWP07S02 5-157 07-0010 Vial Number: 14 14000 13000 12000 11000 0006 8000 7000 6000 Abundance, 10000 File

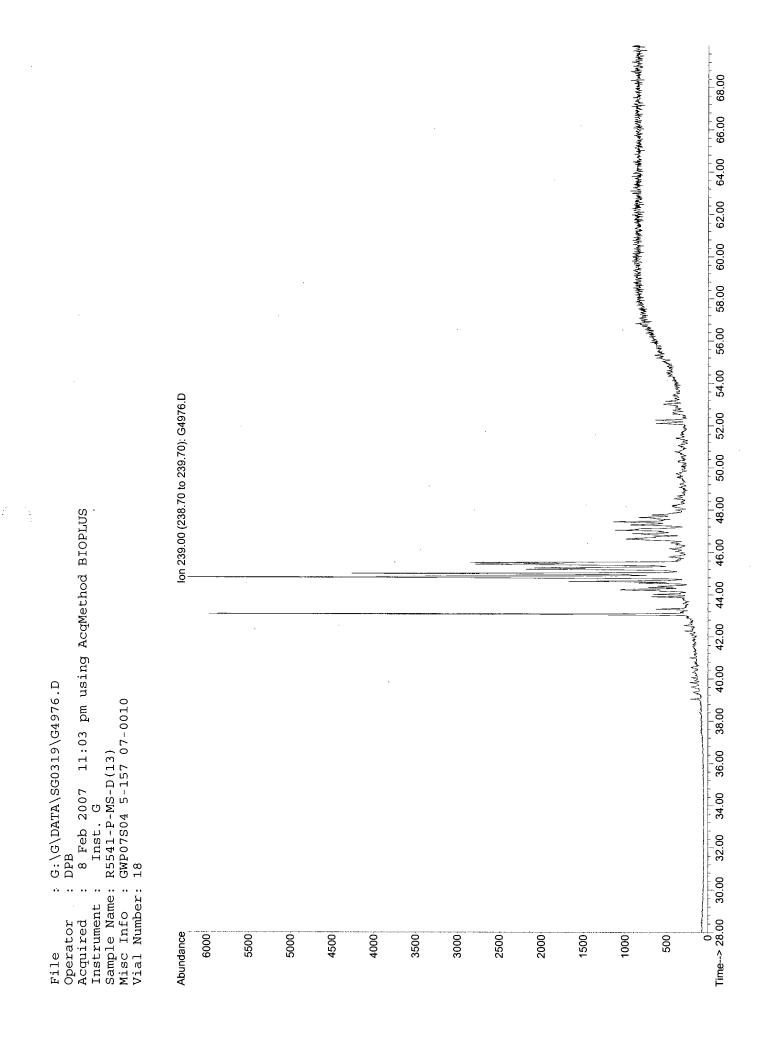
54.00 56.00 58.00 60.00 62.00 64.00 66.00 68.00 52.00 50.00 36.00 38.00 40.00 42.00 44.00 46.00 48.00 NY WWW WW -Time--> 28.00 30.00 32.00 34.00 1 0 1000

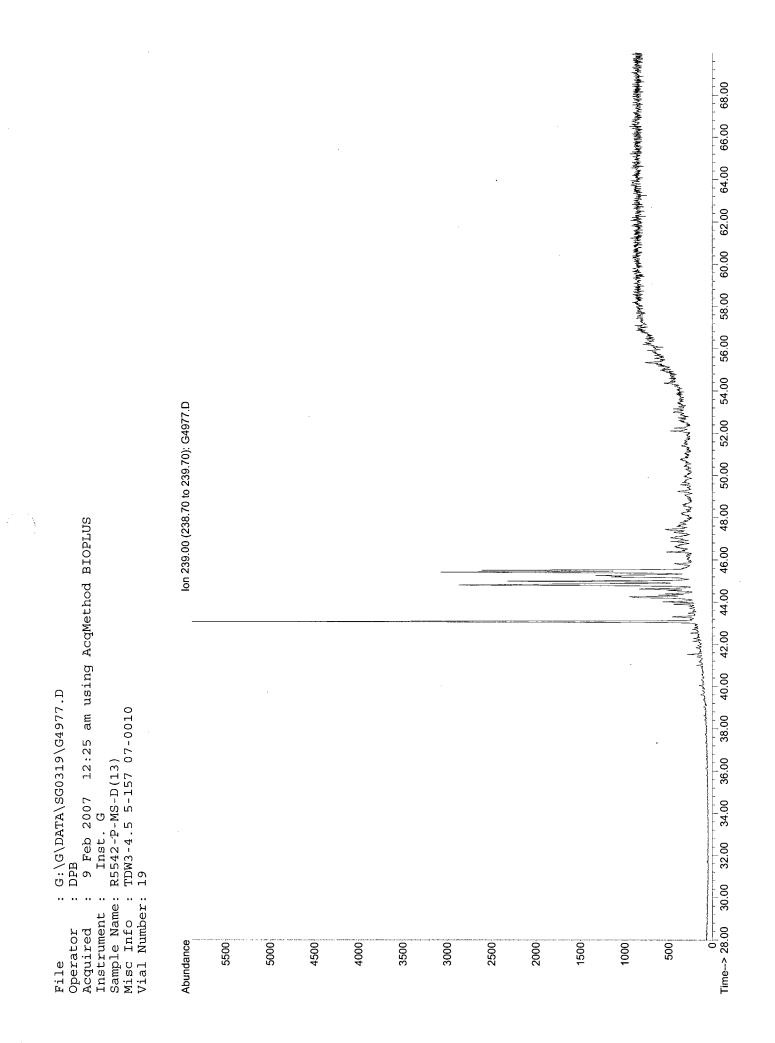
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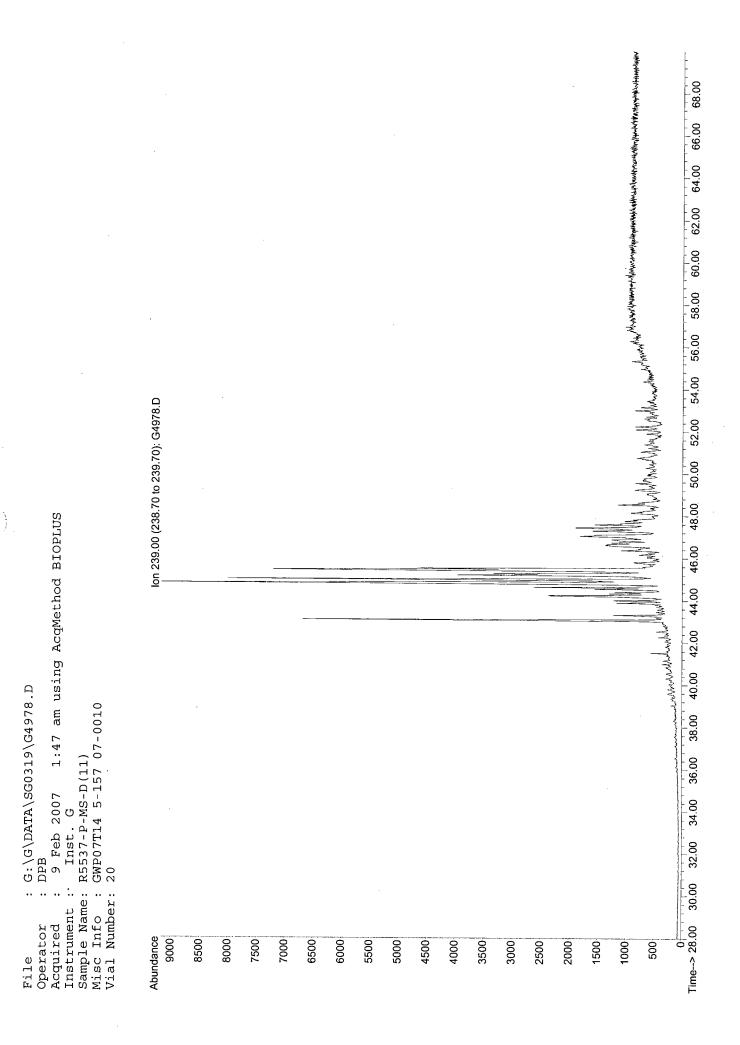
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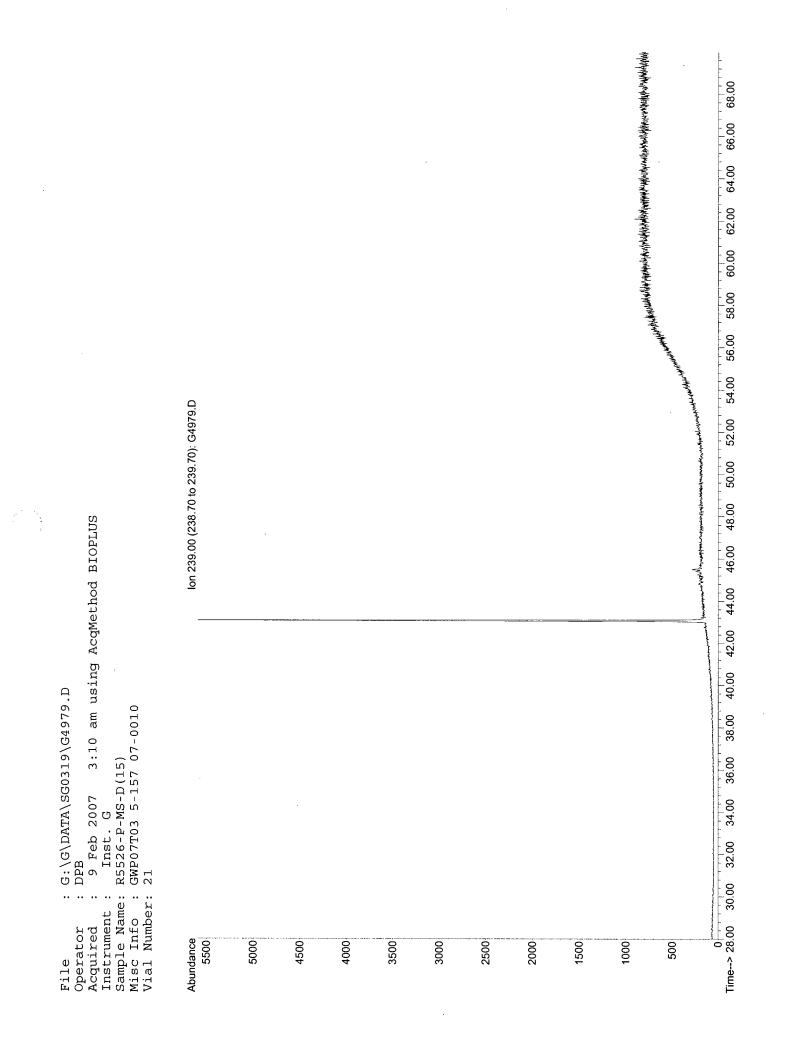
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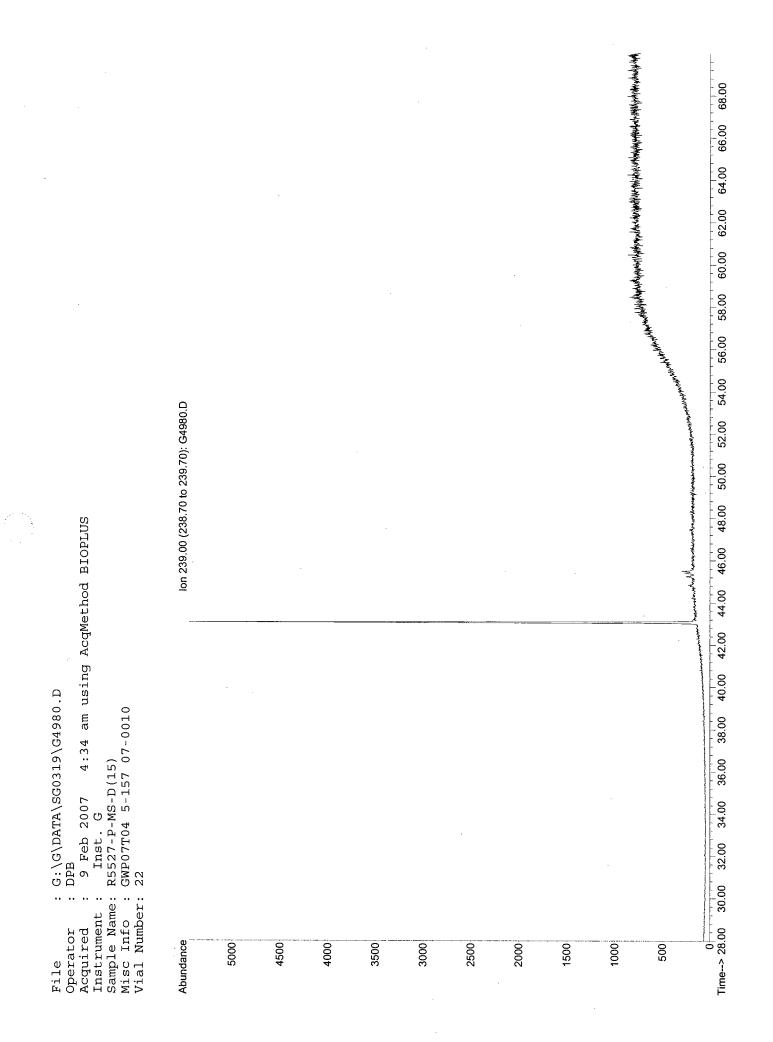
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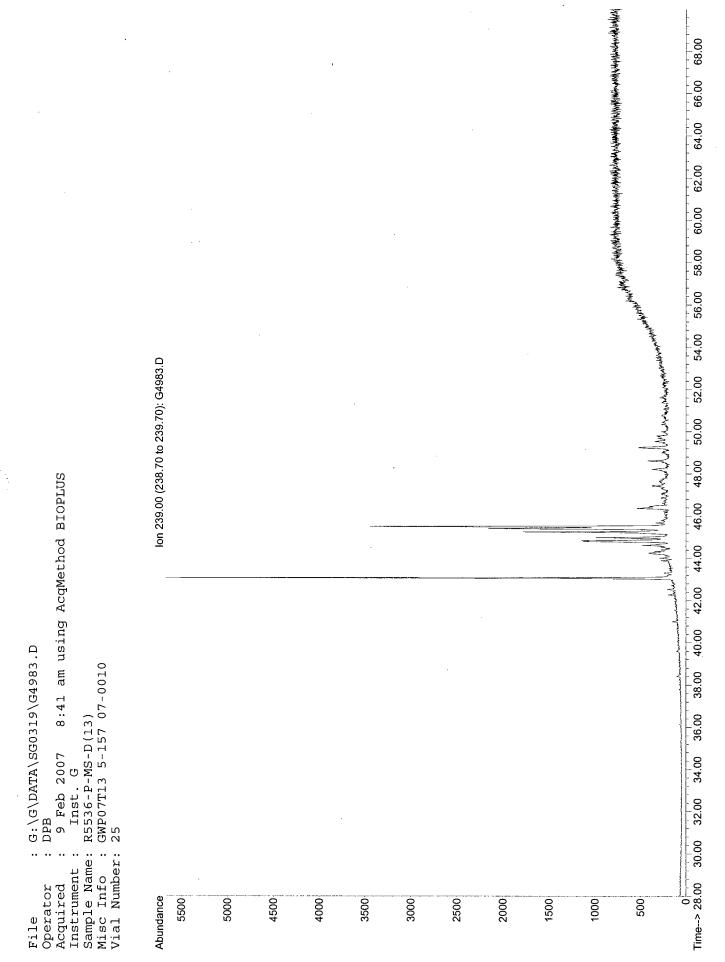




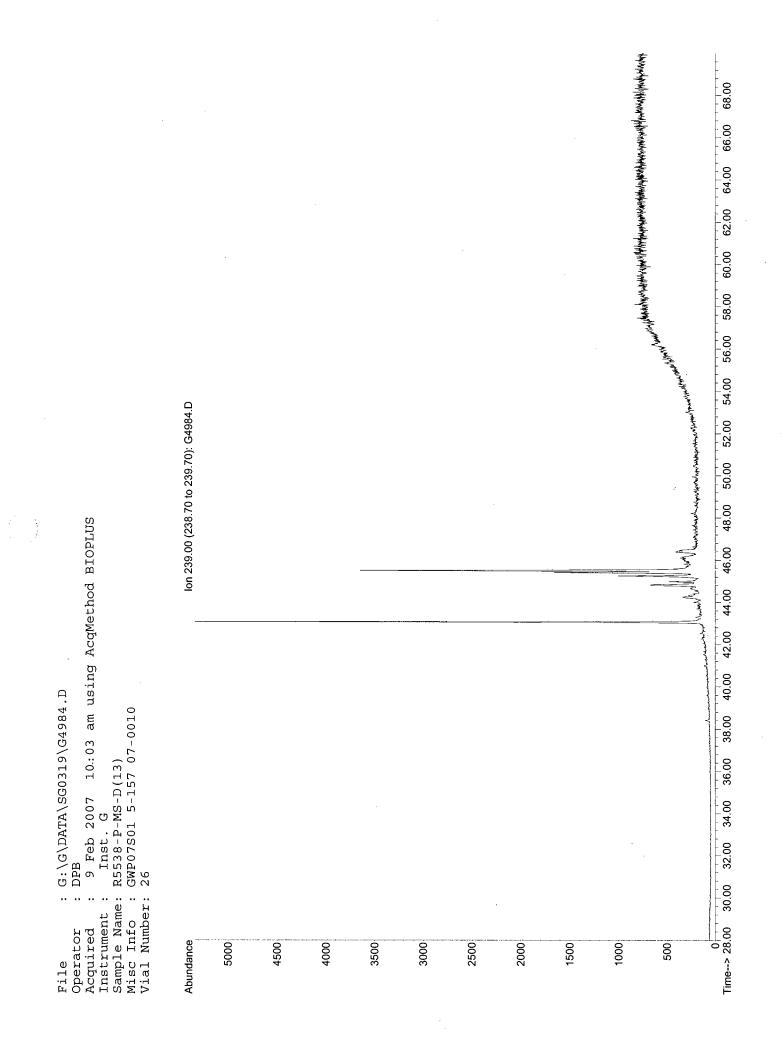


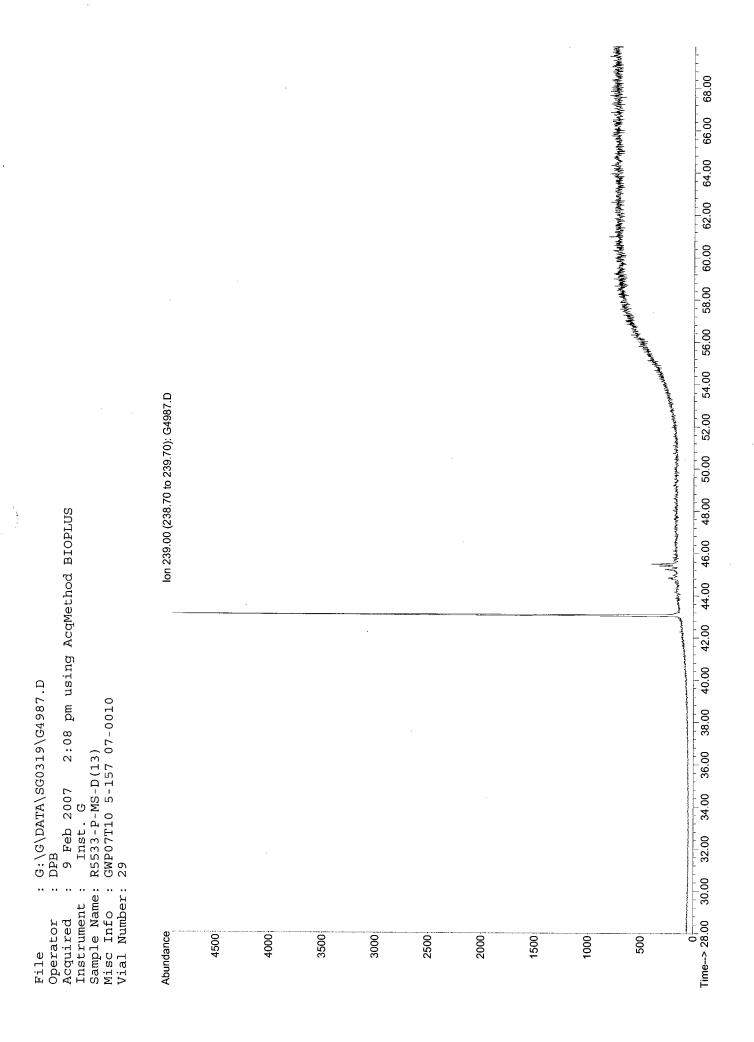


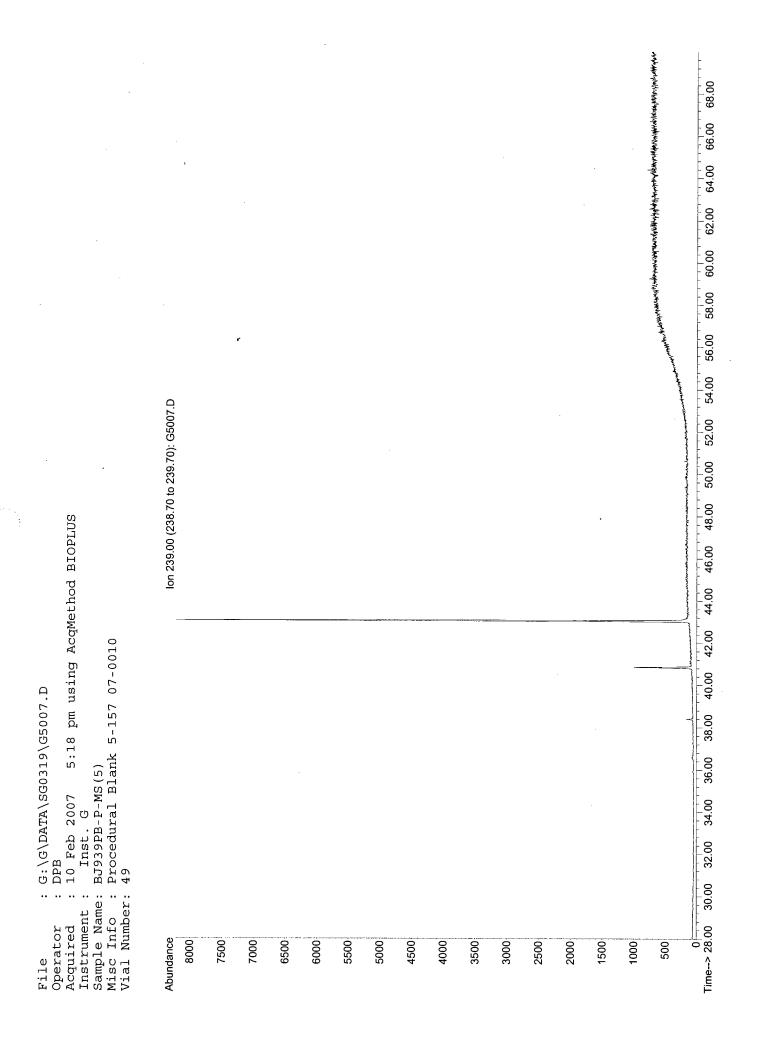


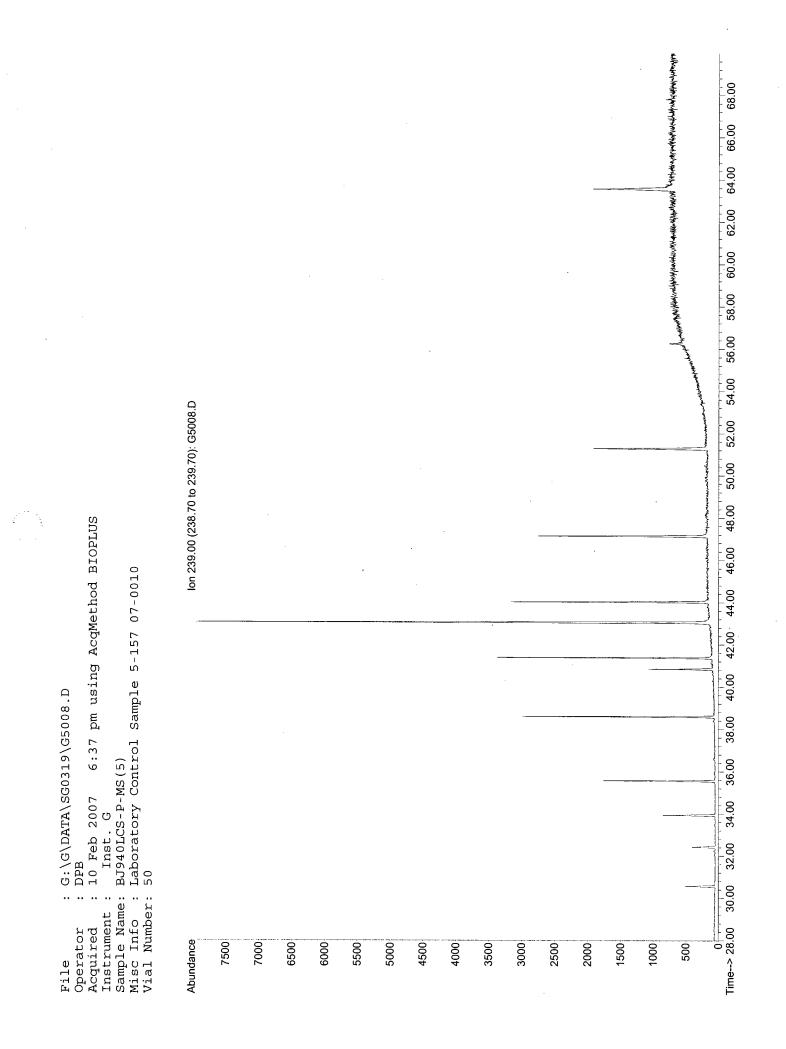


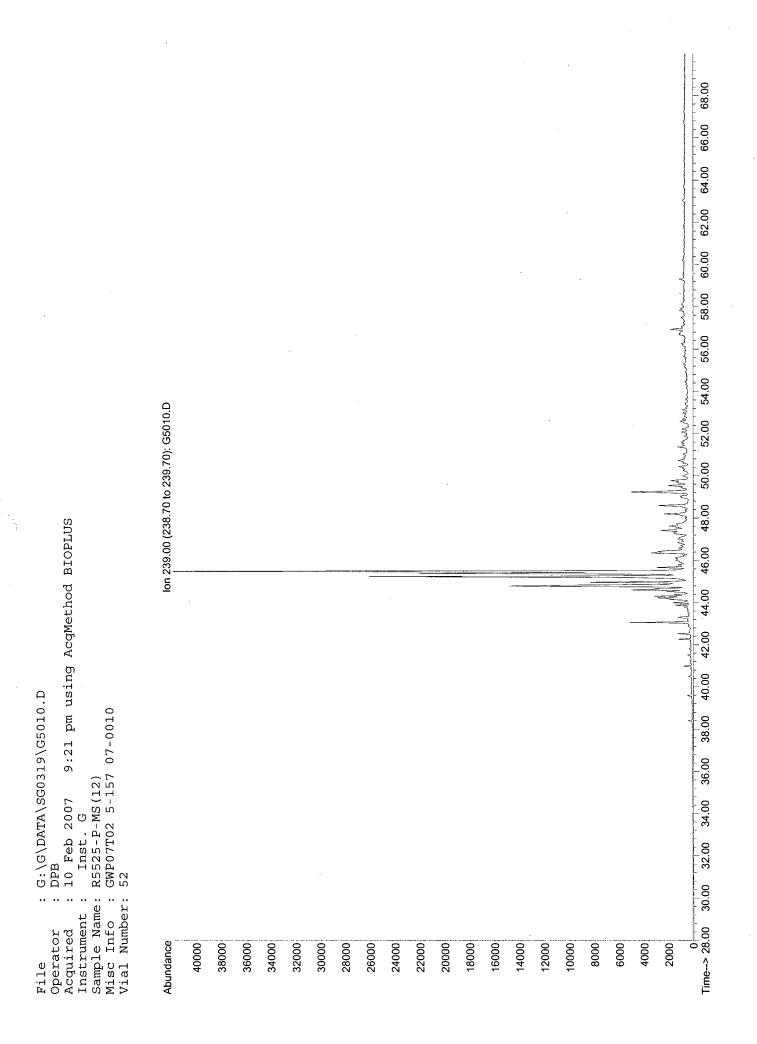
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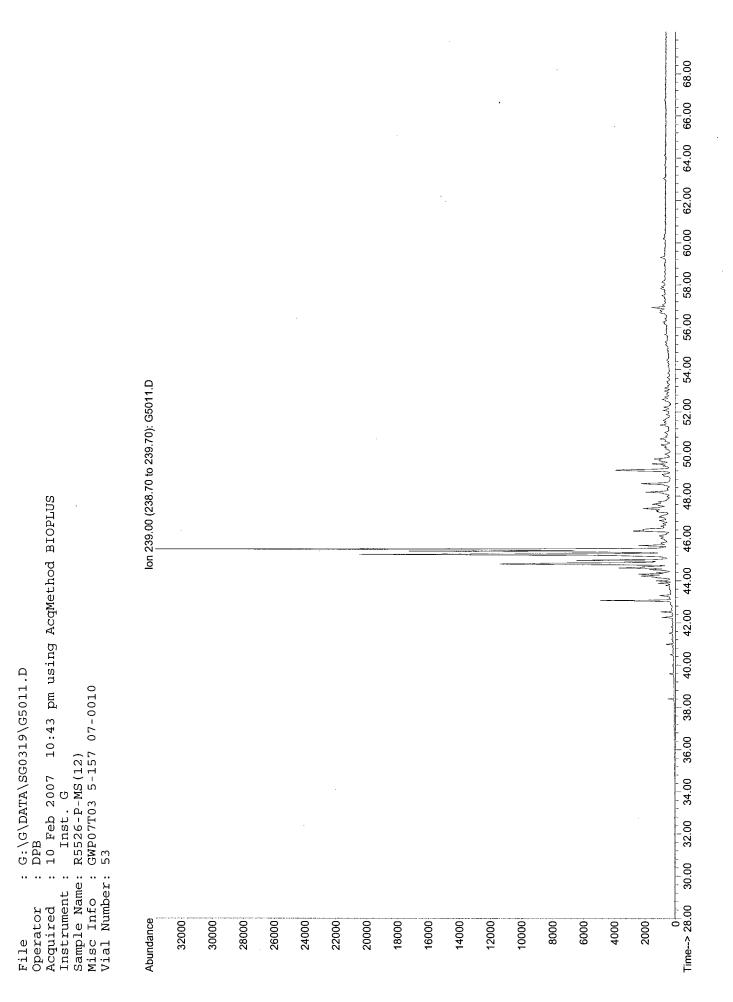




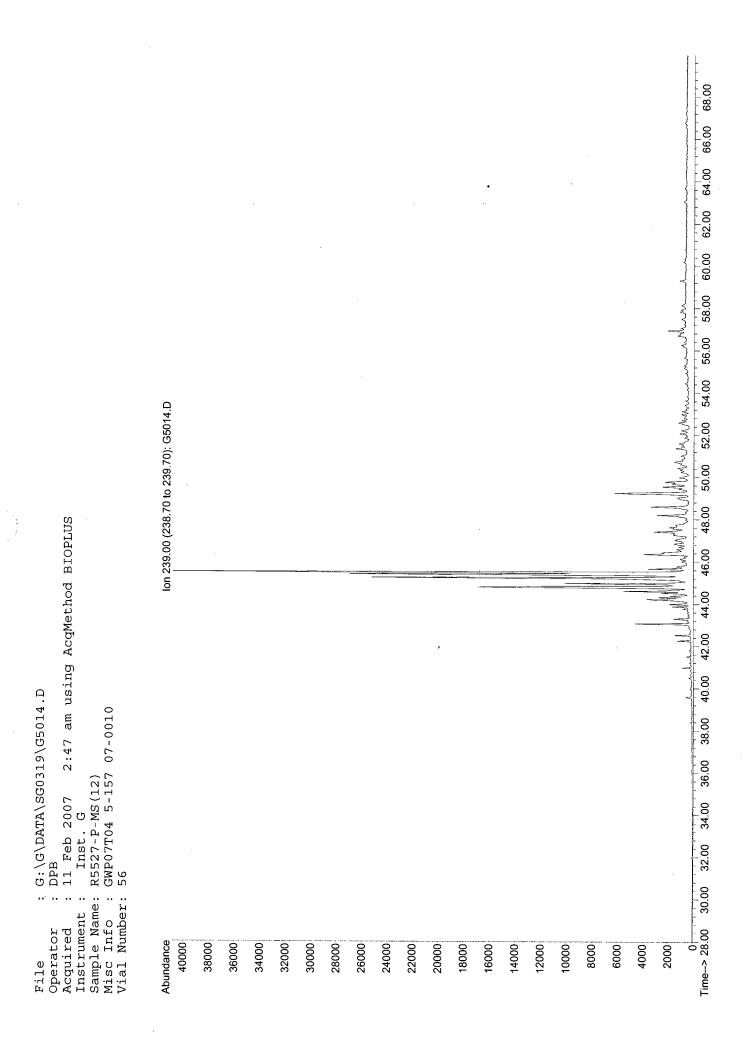


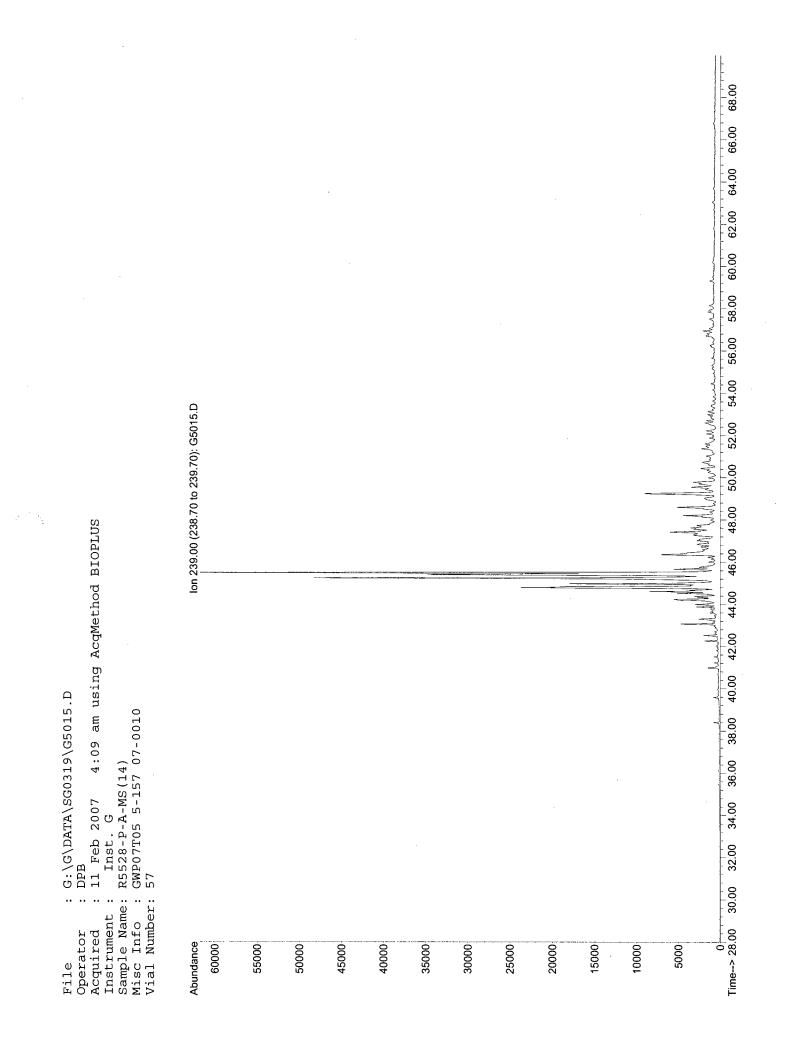


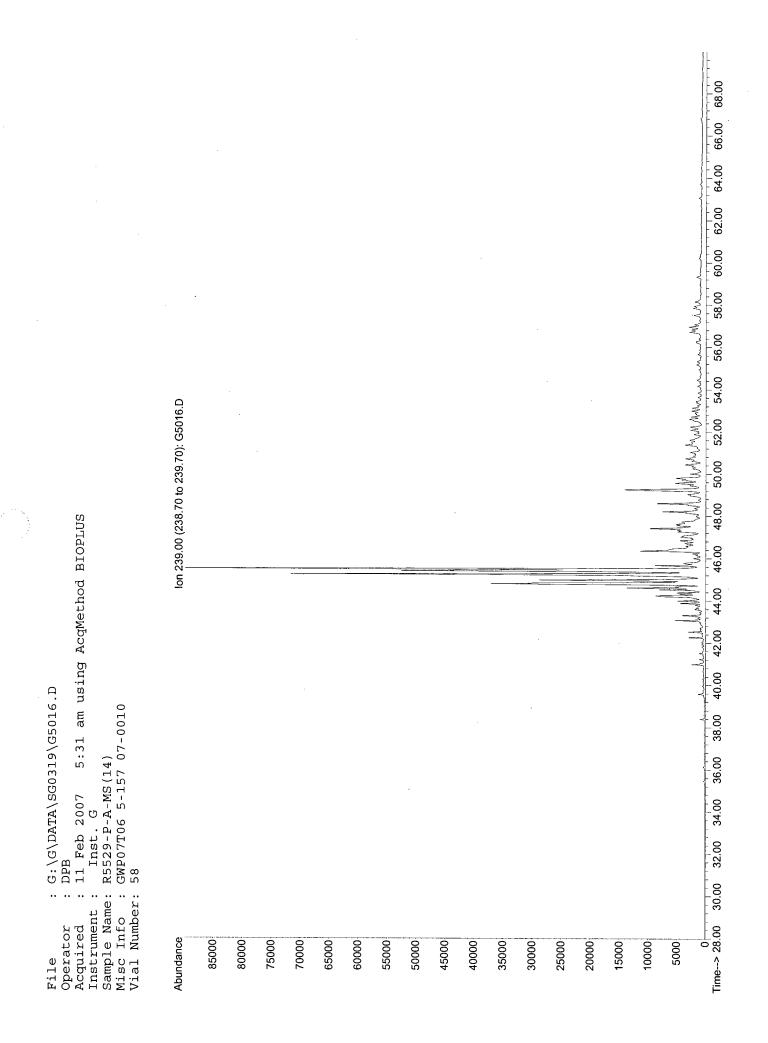


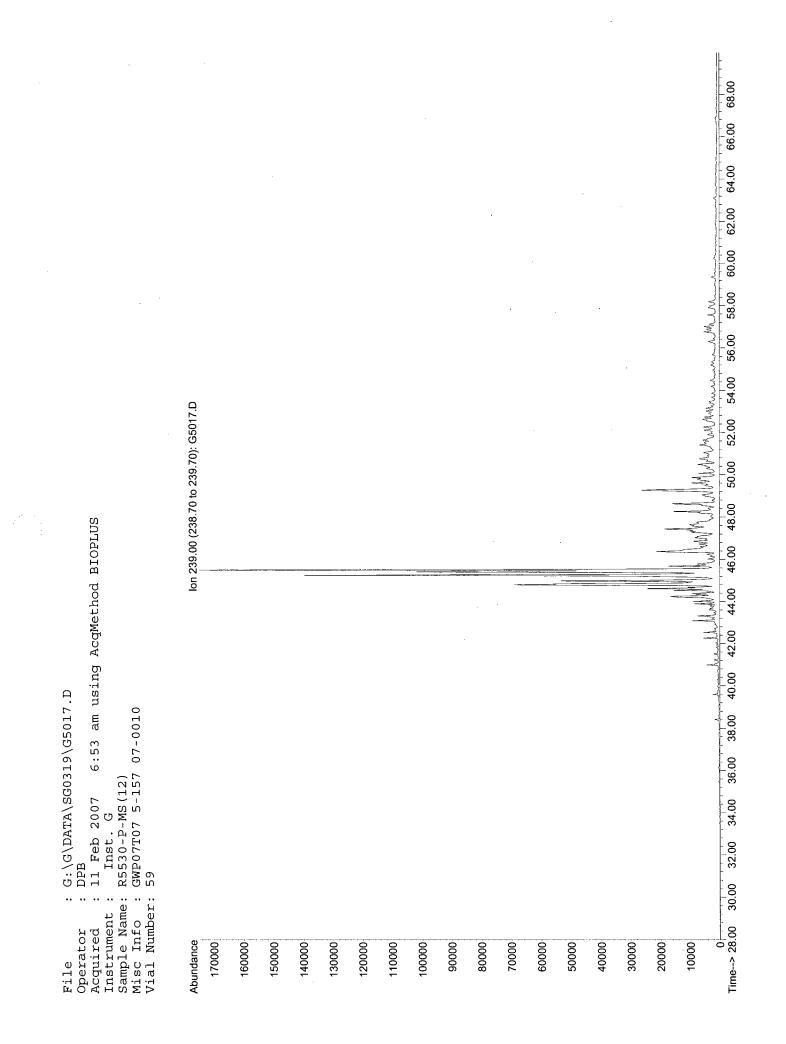


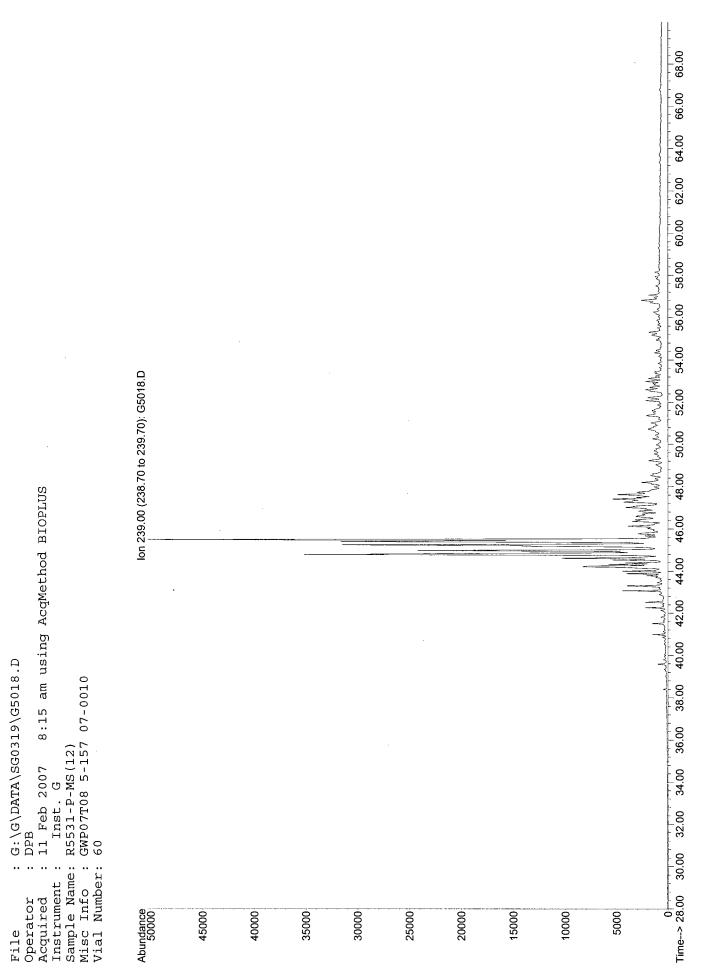
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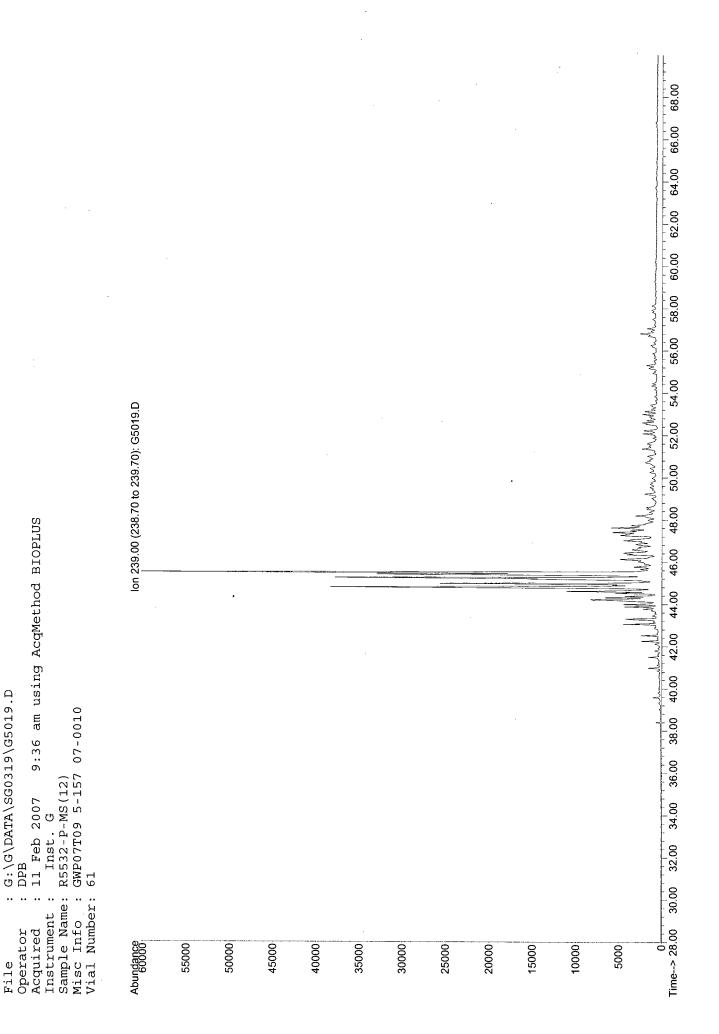


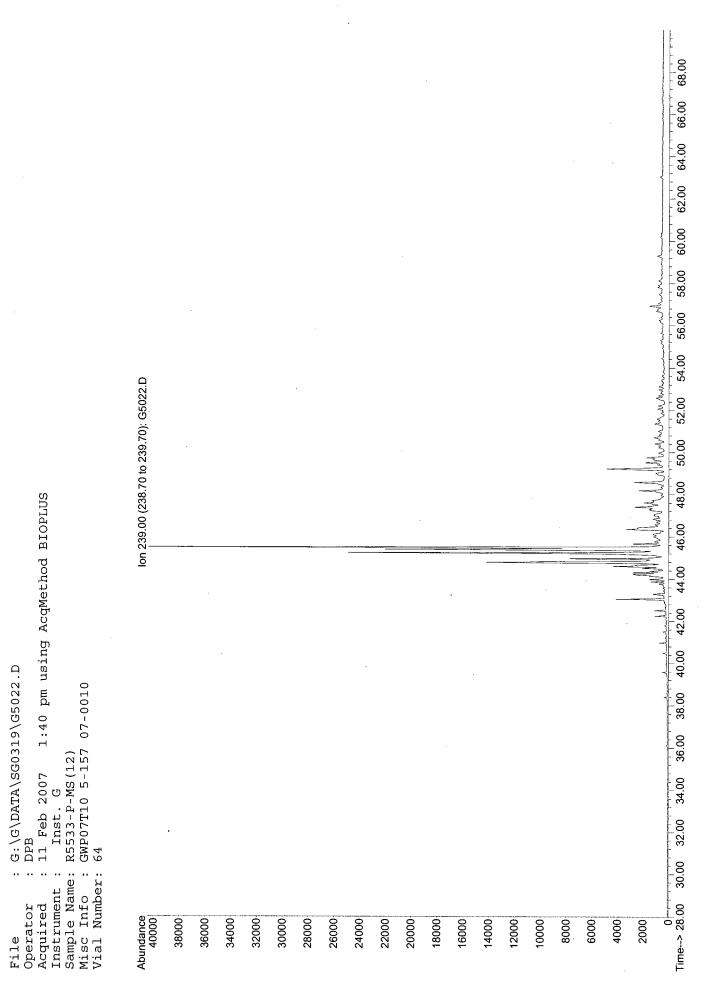


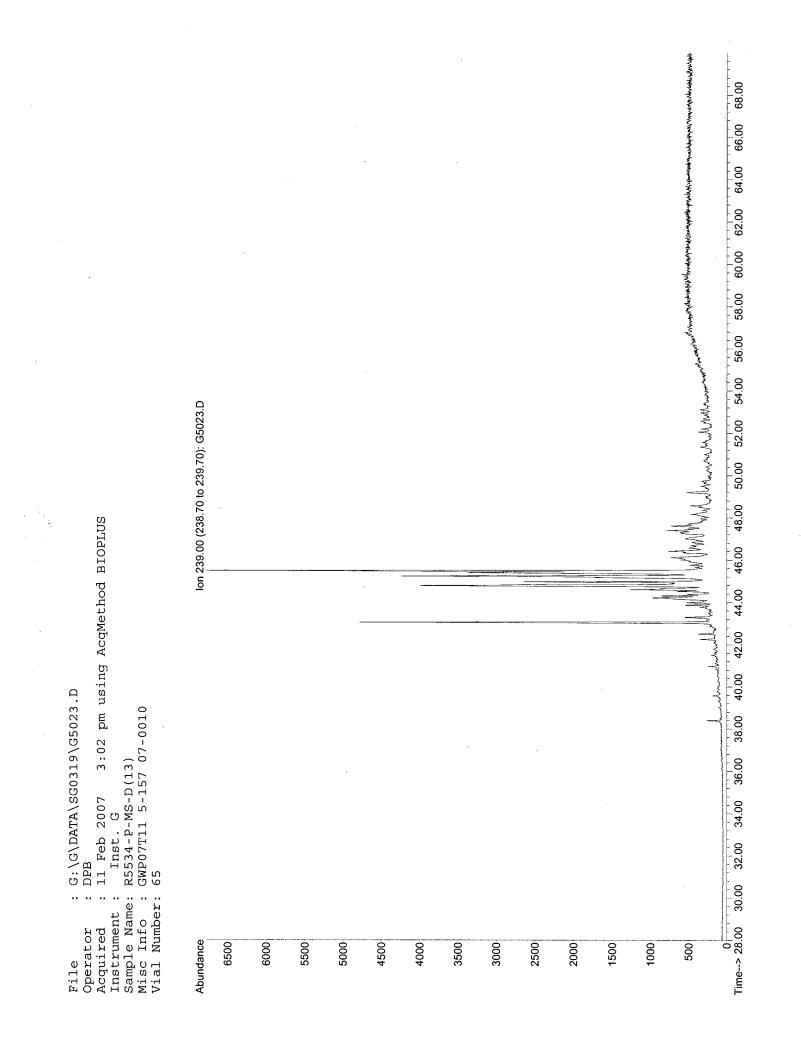


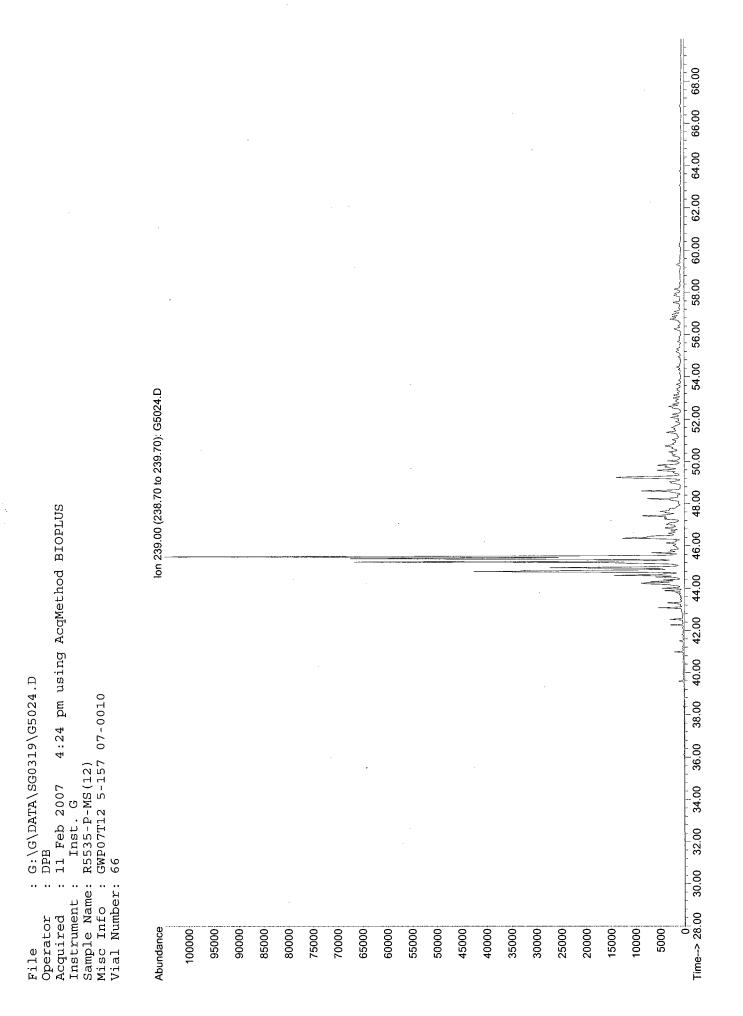
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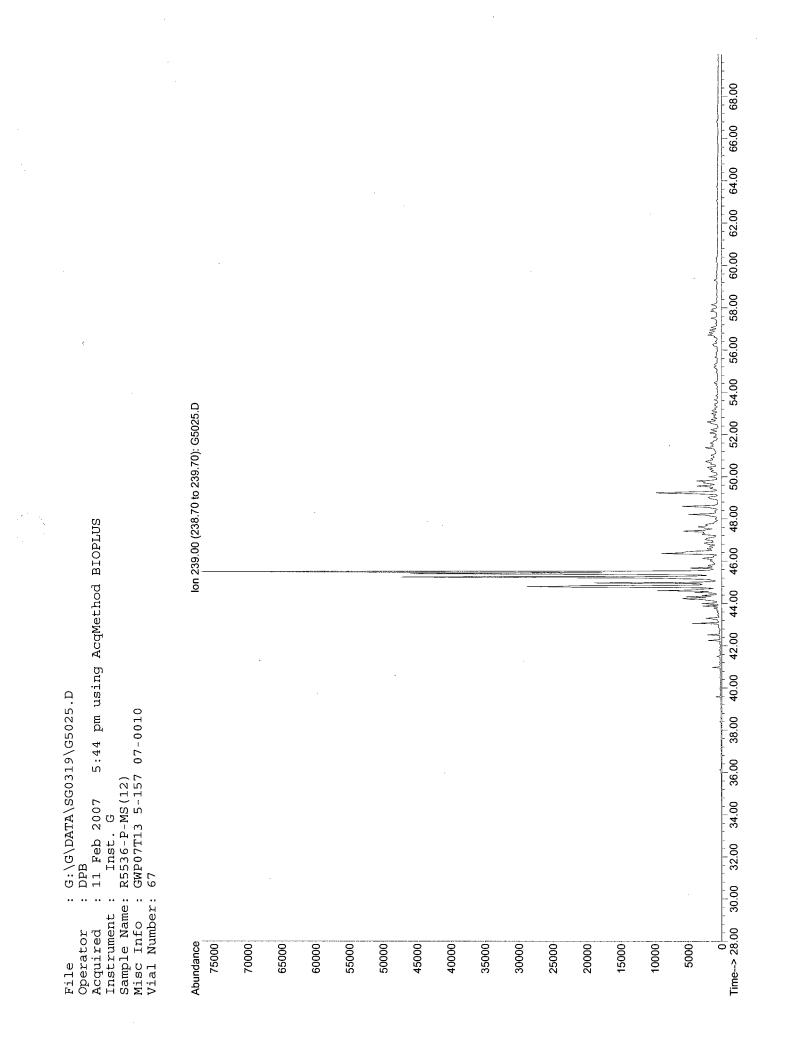


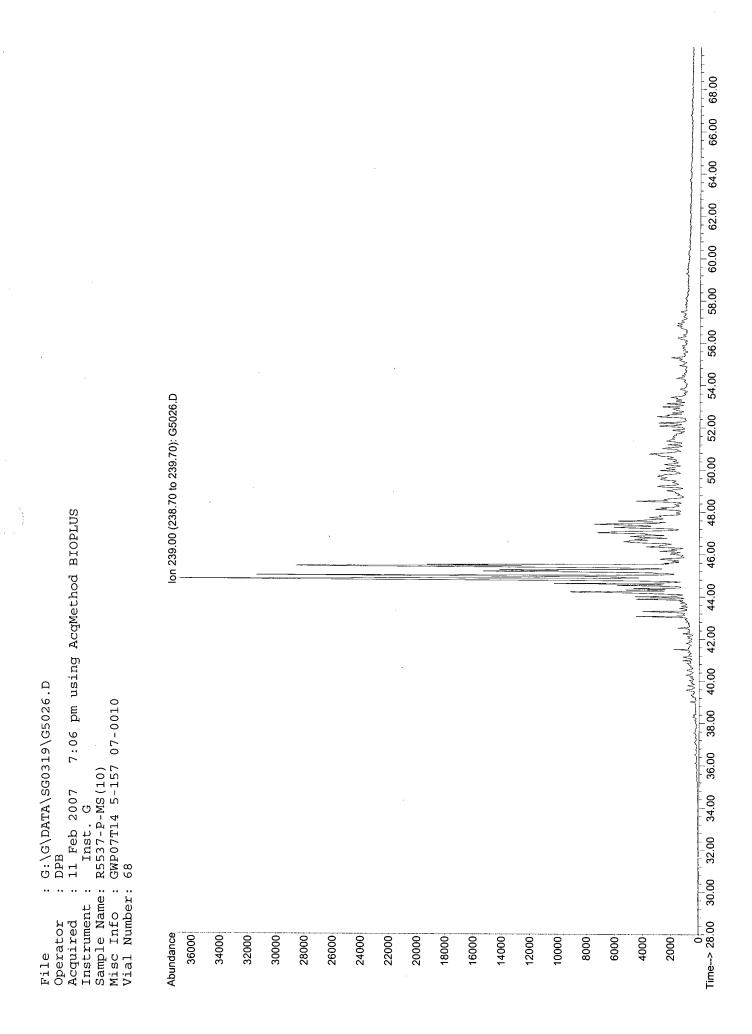


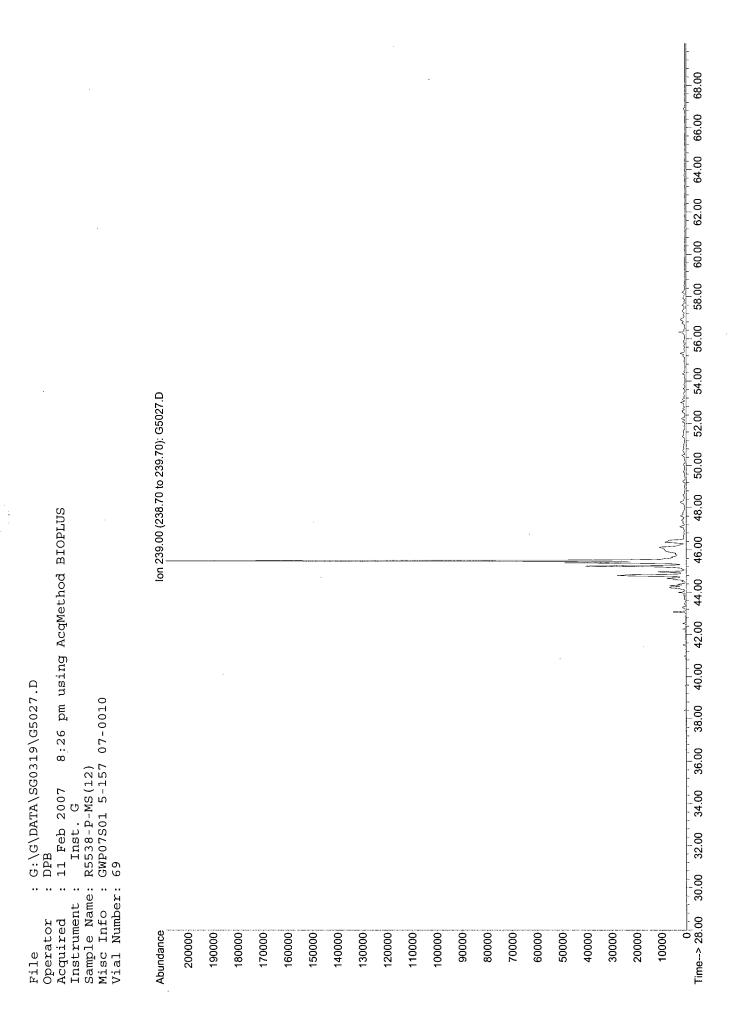


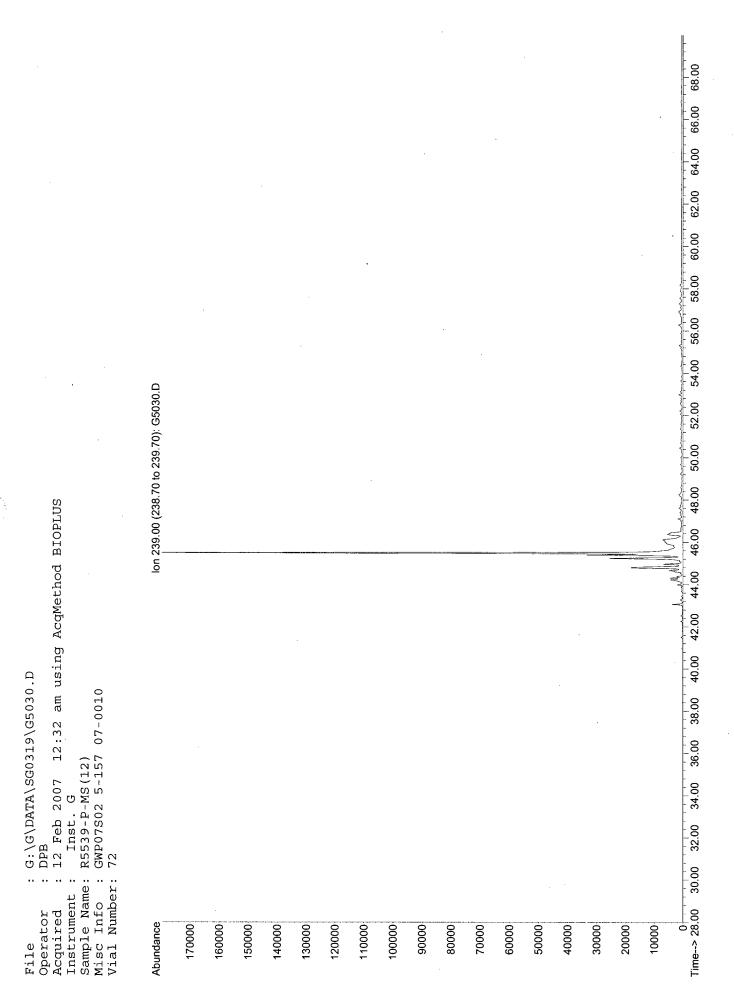


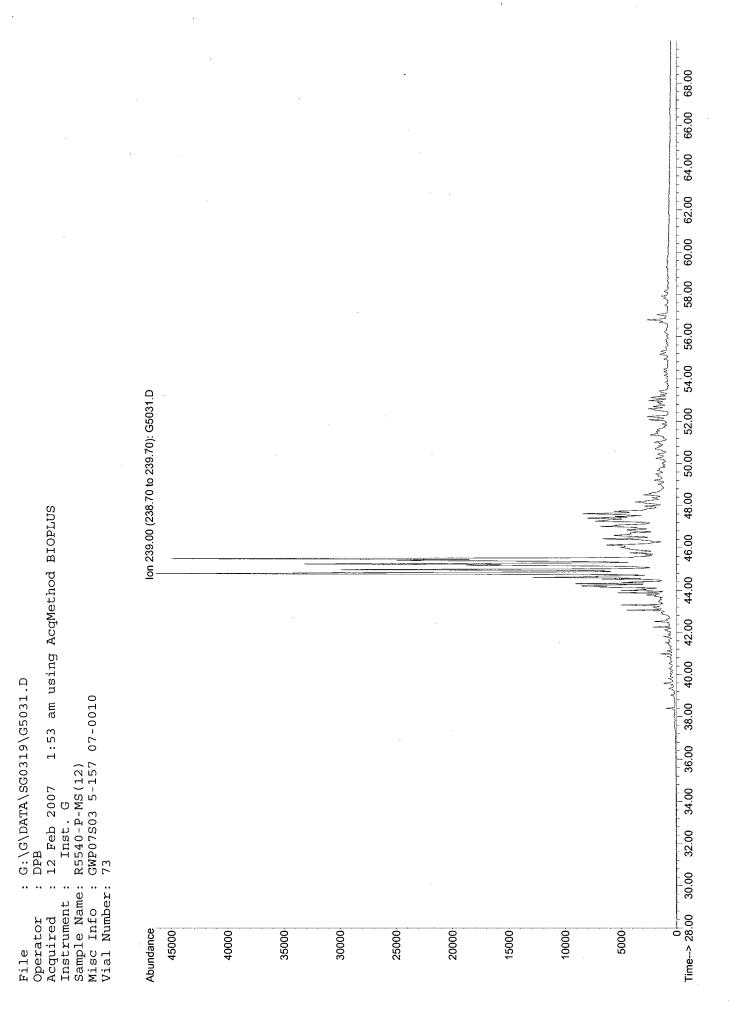
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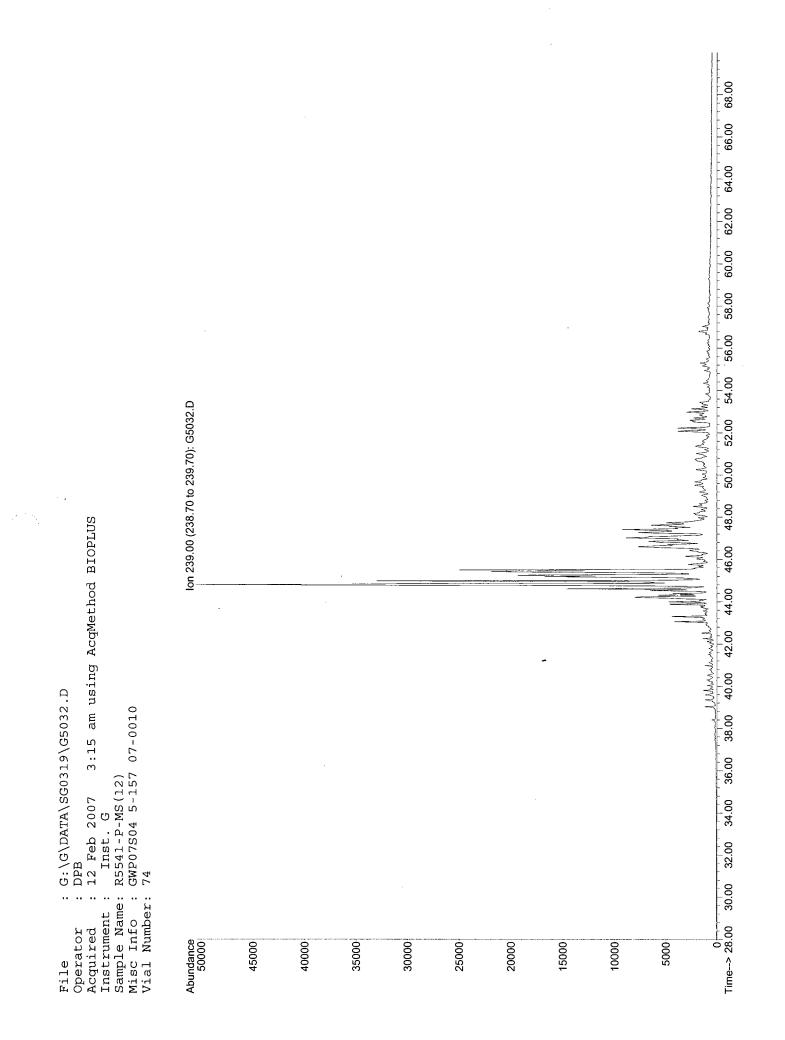


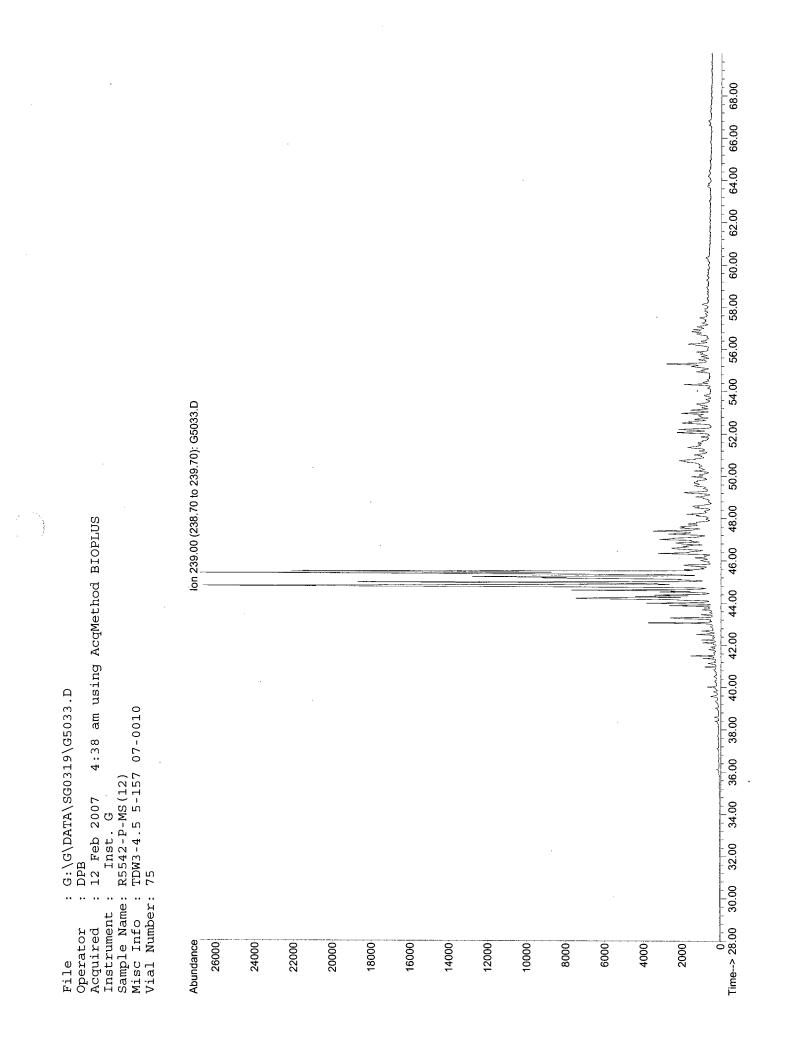


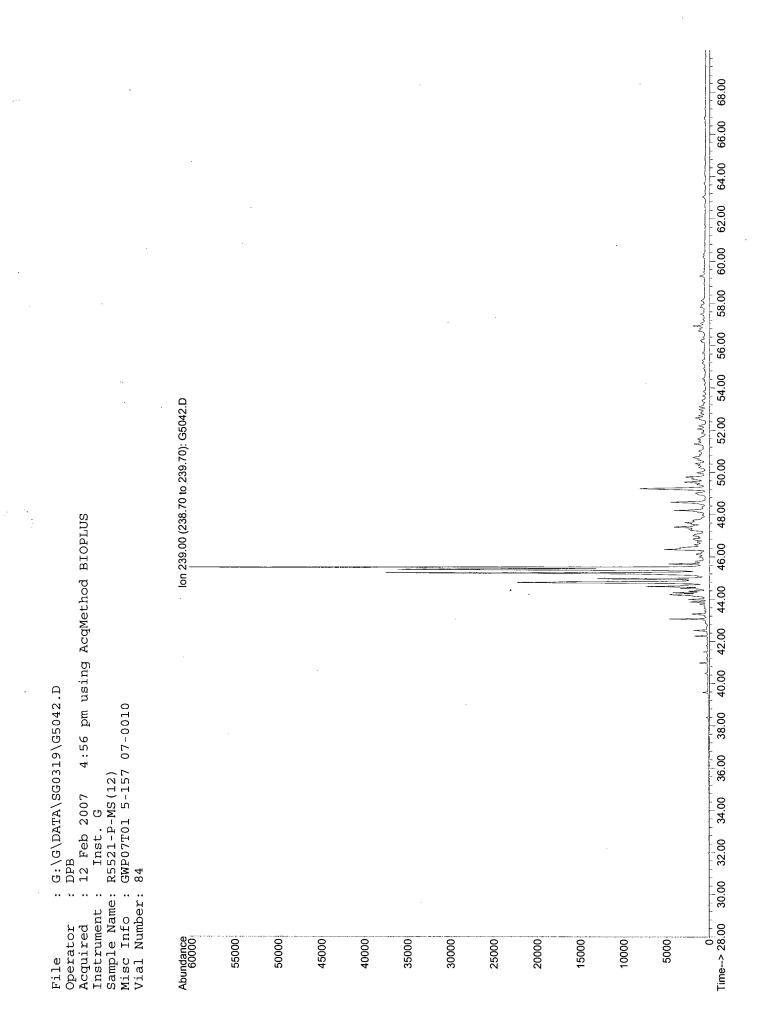












PAH and Biomarker – NAPL QA/QC Summary Batches 07-0011

PROJECT:	Exponent – Gas Works Park
PARAMETER:	Polycyclic Aromatic Hydrocarbons and Biomarkers
LABORATORY:	Battelle, Duxbury, MA
MATRIX:	Non-aqueous phase liquid (NAPL) and Filter/NAPL samples
SAMPLE CUSTODY:	Eighteen tar samples, three NAPLs samples, and 1 soil sample were received at the Battelle
	Duxbury Operations (BDO) Laboratory on 1/16/2007. Upon receipt of samples, the
	temperatures of the coolers were taken and the samples were logged into the laboratory and
	given unique IDs. The temperature of the cooler upon receipt was within the acceptable
	range. Samples were either stored in an access-limited walk-in refrigerator at 4°C until
	sample preparation could begin. The NAPL samples were extracted together in one
	analytical batch, batch 07-0011.

QA/QC DATA QUALITY OBJECTIVES:

	Reference Method	Blank	Surrogate Recovery	LCS/MS Recovery	Control Oil % Diff.	
PAH and petroleum biomarkers	EPA 8270 modified	< 5x MDL	40-120% Recovery	40-120% Recovery MS target spike must be > 5 x background	PD < 30% for 90% of the analytes	_

METHOD:

NAPL samples were prepared for analysis by weighing approximately 50 mg of oil and diluting with 10 mL of hexane. A portion of the extract was removed and spiked with SIS and IS. One extract was submitted for PAH and petroleum biomarker analysis, and the second extract was submitted for SHC and TPH analysis. NAPL sample data is reported on an oil weight basis.

PAH and petroleum biomarkers were measured by gas chromatography-mass spectrometry (GC/MS) in the selected ion mode (SIM) using a modified EPA Method 8270. An initial calibration consisting of target analytes was analyzed prior to analysis to demonstrate the linear range of analysis. Calibration verification was performed at the beginning and end of each 12 hour period in which samples were analyzed. Concentrations of PAH and petroleum biomarkers were calculated versus internal standards. Target PAH were quantified using the average response factors (RF) generated from the initial calibration. The alkyl homolgue PAH series were assigned the RF of the parent PAH, steranes were assigned the RF of cholestane, and triterpanes were assigned the RF of moretane.

Note: the reporting limit for alkylbenzene compounds is orders of magnitude higher than the reporting limit for the rest of the PAH compounds.

HOLDING TIMES: Samples were stored cool at approximately 4°C until extraction.

Samples were prepared for analysis in one analytical batch and analyzed within 40 days of extraction.

Batch ID	Extraction Date	Analysis Date(s)
07-0011	1/30/2007	2/7/2007 - 2/12/2007

PAH and Biomarker – NAPL QA/QC Summary Batches 07-0011

PROCEDURAL BLANK (PB):	A procedural blank (PB) was prepared with each analytical batch. The blank was analyzed to ensure the sample extraction and analysis methods were free of contamination.
	07-0011 – No exceedences noted.
	Comments- None.
LABORATORY CONTROL SAMPLE (LCS):	A laboratory control sample (LCS) was prepared each analytical batch. The percent recoveries of target analytes were calculated to measure data quality in terms of accuracy.
(LCS).	07-0011 – No exceedences noted.
	Comments- None.
SURROGATE RECOVERY:	Five surrogate compounds were added prior to extraction, including d8-naphthalene, d10- acenaphthene, d10-phenanthrene, d12-benzo(a)pyrene, and 5(b)H-cholane. The recovery of the surrogate compound was calculated to measure data quality in terms of accuracy (extraction efficiency).
	07-0011 - 4 exceedences noted.
	Comments –d10-acenaphthtene and 5b(H)-cholane was over-recovered in the following samples: GWP07DW401 and GWP07DW402. These over-recoveries are due to an interfering peak. The exceedences were qualified with an "NME" to indicate the exceedence is an estimate due to matrix interference.
CONTROL OIL:	A control oil (North Slope Crude) was prepared with the analytical batch. The percent difference (PD) between the measured value and the target value was calculated to measure data quality in terms of accuracy.
	07-0011 – No exceedences noted.
	Comments – None.
CALIBRATIONS:	The GC/MS is calibrated with a minimum 5 level curve for all compounds. The percent relative standard deviation (%RSD) between RF for the individual target analytes must be \leq 30%, and the mean RSD of all target analytes must be <15%. Each batch of samples analyzed is bracketed by a continuing calibration verification (CCV) sample, run at a frequency of minimally every 12 hours. The PD between the true value and the CCV should be <25% for individual analytes. Additionally an initial calibration check (ICC) sample is run immediately after each initial calibration. The PD between the ICC and the initial calibration should be <25%.
	07-0011 – No exceedences noted.
	Comments – None.

Battelle The Business of Innovation

Client ID	GWP07DW401	GWP07DW402	GWP07MW9
Battelle ID	R5522-P	R5523-P	R5524-P
Sample Type	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/12/07
Extraction Date	01/30/07	01/30/07	01/30/07
Analysis Date	02/09/07	02/09/07	02/09/07
Analytical Instrument	MS	MS	MS
% Moisture	NA	NA	NA
% Lipid	NA	NA	NA
Matrix	NAPL	NAPL	NAPL
Sample Size	49.50	47.40	45.80
Size Unit-Basis	MG OIL	MG_OIL	MG_OIL
Minimum Reporting Limit	1.38	1.45	1.35
Units	MG/KG_OIL	MG/KG_OIL	MG/KG_OIL
C3-Alkylbenzenes	2138.94	2176.03	872.5
C4-Alkylbenzenes	1395.75	1403.29	1048.71
C5-Alkylbenzenes	333.04	331.6	405.07
C6-Alkylbenzenes	122.64 J	125.34 J	443.22
Benzo(b)thiophene	2816.79	2829.81	198.62
C1-benzo(b)thiophenes	2034.58	2036.28	493.55
C1-benzo(b)(hiophenes C2-benzo(b)thiophenes	2034.58 1687.46	1685.17	493.55 621.88
C2-benzo(b)(hiophenes C3-benzo(b)thiophenes	728.13	717.84	533.11
C3-benzo(b)(hiophenes C4-benzo(b)thiophenes	211.42	211.15	306.86
Naphthalene	80881 D	83156.26 D	11835.73 D
•	32020.01	32249.35	9666.3
C1-Naphthalenes		17857.55	7679.23
C2-Naphthalenes	17923.86 6609.39		
C3-Naphthalenes		6549.99	4057.14
C4-Naphthalenes	1605.29	1593.48	1767.19
C1-Biphenyls + Dibenzofuran	7569.69	7549.04	917.96
Biphenyl	4858.56 3603.81	4845.03	626.06
C2-Biphenyls + C1-Dibenzofurans		3590.56	1299.63
Acenaphthylene	5833.45	5845.03	546.25
Acenaphthene	7215.98	7207.79	1882.87
Dibenzofuran	4859.52	4833.66	216.65
Fluorene	7591.66 D	7806.98 D	1491.6
C1-Fluorenes	2673.78	2621.67	1231.7
C2-Fluorenes	1131.96	1102.6	883.48
C3-Fluorenes C4-Fluorenes	369.59	373.58	445.92
Anthracene	312.01 7275.9	311.66 7158.32	429.99 877.01
Phenanthrene	22250.58 D	22825.88 D	3626.27
C1-Phenanthrenes/Anthracenes	8916.84	8704.98	2830.13
C2-Phenanthrenes/Anthracenes	3174.56	3075.38	1432.77
C3-Phenanthrenes/Anthracenes	821.77	783.37	566.25
C4-Phenanthrenes/Anthracenes	184.59	186.81	149.46
Retene	187.75	182.9	82.04
Dibenzothiophene	1693.56	1658.84	339.8
•	1123.6	1096.14	475.18
C1-Dibenzothiophenes C2-Dibenzothiophenes	592.49	569.64	399.12
C3-Dibenzothiophenes C4-Dibenzothiophenes	218.44 59.24	218.21	221.59 78.75
Fluoranthene	7751.27 D	56.66 7953.79 D	969.93
Pyrene	8050.52 D	8352.58 D	1497.93
C1-Fluoranthenes/Pyrenes			
C2-Fluoranthenes/Pyrenes	5167.6 1387.61	5008.93 1373.09	1405.69 541.28
C2-Fluoranthenes/Pyrenes			202.56
C4-Fluoranthenes/Pyrenes	427.01	417.1 142.48	71.92
C0-Benzo(b)naphthothiophenes	146.9 271.49	142.48	47.51
C0-Benzo(b)naphthothiophenes	271.49 221.3	262.16 231.36	76.72
C1-Benzo(b)naphthothiophenes	92.05	88.89	46.1
C2-Benzo(b)naphthothiophenes	92.05 52.73		26.65
		53.85	
C4-Benzo(b)naphthothiophenes	13.7	16.28	8.88 547 84
Benzo(a)anthracene	3227.37	3279.55 3054.96	547.84
Chrysene	3000.92	5054.90	535.57

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	GWP07DW401	GWP07DW402	GWP07MW9
Battelle ID	R5522-P	R5523-P	R5524-P
Sample Type	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/12/07
Extraction Date	01/30/07	01/30/07	01/30/07
Analysis Date	02/09/07	02/09/07	02/09/07
Analytical Instrument	MS	MS	MS
% Moisture	NA	NA	NA
% Lipid	NA	NA	NA
Matrix	NAPL	NAPL	NAPL
Sample Size	49.50	47.40	45.80
Size Unit-Basis	MG OIL	MG OIL	MG OIL
Minimum Reporting Limit	1.38	1.45	1.35
Units	MG/KG OIL	MG/KG OIL	MG/KG OIL
	_		
C1-Chrysenes	1471.93	1508.6	414.23
C2-Chrysenes	507.39	517.36	176.04
C3-Chrysenes	282.15	300.06	87.51
C4-Chrysenes	43.48	46.28	12.88
Benzo(b)fluoranthene	1546.11	1593.49	187.89
Benzo(k)fluoranthene	2222.36	2223.93	276.86
Benzo(e)pyrene	1581.81	1618.3	213.72
Benzo(a)pyrene	3163.62	3212.24	454.19
Perylene	705.74	721.08	73.34
Indeno(1,2,3-cd)pyrene	1555.21	1595.97	170.51
Dibenz(a,h)anthracene	326.56	337.24	47.31
Benzo(g,h,i)perylene	1437.13	1473.18	165.02
Total PAH	263741.41	266964.93	61306.75
Surrogate Recoveries (%)			
Naphthalene-d8	102	100	91
Acenaphthene-d10	132 NME	128 NME	99
Phenanthrene-d10	96	93	89
Benzo(a)pyrene-d12	104	107	112
		101	

The Business of Innovation

Client ID	Procedural Blank	
Battelle ID	BJ941PB-P	
Sample Type	PB	
Collection Date	01/30/07	
Extraction Date	01/30/07	
Analysis Date	02/12/07	
Analytical Instrument	MS	
% Moisture	NA	
% Lipid	NA	
Matrix	NAPL	
Sample Size	5.00	
Size Unit-Basis	MG_OIL	
Minimum reporting Limit	1.3	
Units	MG/KG_OIL	
C3-Alkylbenzenes	U	
C4-Alkylbenzenes	U	
C5-Alkylbenzenes	U	
C6-Alkylbenzenes	U	
Benzo(b)thiophene	U	
C1-benzo(b)thiophenes	U	
C2-benzo(b)thiophenes	U	
C3-benzo(b)thiophenes	U	
C4-benzo(b)thiophenes	U	
Naphthalene	0.44 J	
C1-Naphthalenes	U	
C2-Naphthalenes	U	
C3-Naphthalenes	U	
C4-Naphthalenes	U	
C1-Biphenyls + Dibenzofuran	U	
Biphenyl	U	
C2-Biphenyls + C1-Dibenzofurans	U	
Acenaphthylene	U	
Acenaphthene	U	
Dibenzofuran	U	
Fluorene	U	
C1-Fluorenes	U	
C2-Fluorenes	U	
C3-Fluorenes	U	
C4-Fluorenes	U	
Anthracene	U	
Phenanthrene	0.14 J	
C1-Phenanthrenes/Anthracenes	U	
C2-Phenanthrenes/Anthracenes	U	
C3-Phenanthrenes/Anthracenes	U	
C4-Phenanthrenes/Anthracenes	U U	
Retene	U	
Dibenzothiophene C1-Dibenzothiophenes	U	
C1-Dibenzothiophenes	U	
C3-Dibenzothiophenes		
C3-Dibenzothiophenes	U U	
Fluoranthene	0.07 J	
Pyrene	0.07 J 0.06 J	
C1-Fluoranthenes/Pyrenes	0.00 J U	
C2-Fluoranthenes/Pyrenes	U	
C3-Fluoranthenes/Pyrenes	U	
C4-Fluoranthenes/Pyrenes	U	
C0-Benzo(b)naphthothiophenes	U	
C1-Benzo(b)naphthothiophenes	U	
C2-Benzo(b)naphthothiophenes	U	
C3-Benzo(b)naphthothiophenes	U	
C4-Benzo(b)naphthothiophenes	U	
Benzo(a)anthracene	U	
Chrysene	U	
- ,	0	

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	Procedural Blank	
Battelle ID	BJ941PB-P	
Sample Type	PB	
Collection Date	01/30/07	
Extraction Date	01/30/07	
Analysis Date	02/12/07	
Analytical Instrument	MS	
% Moisture	NA	
% Lipid	NA	
Matrix	NAPL	
Sample Size	5.00	
Size Unit-Basis	MG_OIL	
Minimum reporting Limit	1.3	
Units	MG/KG_OIL	
C1-Chrysenes	U	
C2-Chrysenes	U	
C3-Chrysenes	U	
C4-Chrysenes	U	
Benzo(b)fluoranthene	U	
Benzo(k)fluoranthene	U	
Benzo(e)pyrene	U	
Benzo(a)pyrene	U	
Perylene	U	
Indeno(1,2,3-cd)pyrene	U	
Dibenz(a,h)anthracene	U	
Benzo(g,h,i)perylene	U	
Total PAH	0.71 J	
Surrogate Recoveries (%)		
Naphthalene-d8	118	
Acenaphthene-d10	109	
Phenanthrene-d10	105	
Benzo(a)pyrene-d12	93	
Denze(a)pyrene-urz	55	

The Business of Innovation

	Laboratory Control			
Client ID	Sample			
Battelle ID	BJ942LCS-P			
Sample Type	LCS			
Collection Date	01/30/07			
Extraction Date	01/30/07			
Analysis Date	02/12/07			
Analytical Instrument	MS			
% Moisture	NA			
% Lipid	NA			
Matrix	NAPL			
Sample Size	NA			
Size Unit-Basis	NA			
Minimum reporting Limit Units	7.01 NG	Target %	Recovery Qualifier	
	110	Target /	Recovery Qualifier	
C3-Alkylbenzenes	L			
C4-Alkylbenzenes	L			
C5-Alkylbenzenes	L			
C6-Alkylbenzenes				
Benzo(b)thiophene	1160.46	1000.70	116	
C1-benzo(b)thiophenes	L			
C2-benzo(b)thiophenes	L			
C3-benzo(b)thiophenes C4-benzo(b)thiophenes				
Naphthalene	1182.31	1000.20	118	
C1-Naphthalenes	1102.01		110	
C2-Naphthalenes	l			
C3-Naphthalenes	l			
C4-Naphthalenes	ι	J		
C1-Biphenyls + Dibenzofuran	ι	J		
Biphenyl	1170.82	1001.70	117	
C2-Biphenyls + C1-Dibenzofurans	L	J		
Acenaphthylene	1139.35	1000.90	114	
Acenaphthene	1186.2	1000.65	119	
Dibenzofuran	1172.15	1001.90	117	
Fluorene	1152.67	. 1000.55	115	
C1-Fluorenes	l			
C2-Fluorenes	l			
C3-Fluorenes C4-Fluorenes	L			
Anthracene	1124.71	1000.15	112	
Phenanthrene	1181.82	1000.50	118	
C1-Phenanthrenes/Anthracenes	L 1101.02		110	
C2-Phenanthrenes/Anthracenes	l			
C3-Phenanthrenes/Anthracenes	l			
C4-Phenanthrenes/Anthracenes	Ĺ			
Retene	L	J		
Dibenzothiophene	1155.88	1004.60	115	
C1-Dibenzothiophenes	ι			
C2-Dibenzothiophenes	L			
C3-Dibenzothiophenes	L			
C4-Dibenzothiophenes	L			
Fluoranthene	1128.95	1000.50	113	
Pyrene	1092.09	1000.35	109	
C1-Fluoranthenes/Pyrenes	L			
C2-Fluoranthenes/Pyrenes	Ĺ			
C3-Fluoranthenes/Pyrenes C4-Fluoranthenes/Pyrenes	L			
C0-Benzo(b)naphthothiophenes	L L			
C1-Benzo(b)naphthothiophenes				
C2-Benzo(b)naphthothiophenes	l			
C3-Benzo(b)naphthothiophenes	l			
C4-Benzo(b)naphthothiophenes	l			
Benzo(a)anthracene	909.06	1000.25	91	
• •				

The Business of Innovation

Client ID Battelle ID	Laboratory Control Sample BJ942LCS-P				
Sample Type Collection Date	LCS 01/30/07				
Extraction Date	01/30/07				
Analysis Date	02/12/07				
Analytical Instrument	02/12/07 MS				
% Moisture	NA				
% Lipid	NA				
Matrix	NAPL				
Sample Size	NA				
Size Unit-Basis	NA				
Minimum reporting Limit	7.01				
Units	NG		Target %	Recovery	Qualifier
			Ť	· · ·	
Chrysene	1014.88		1000.40	101	
C1-Chrysenes		U			
C2-Chrysenes		U			
C3-Chrysenes		U			
C4-Chrysenes		U			
Benzo(b)fluoranthene	916.03		1000.85	92	
Benzo(k)fluoranthene	1033.98		1000.55	103	
Benzo(e)pyrene	971.81		1002.50	97	
Benzo(a)pyrene	983.44		1000.80	98	
Perylene	915.48		1002.05	91	
Indeno(1,2,3-cd)pyrene	910.17		1000.50	91 00	
Dibenz(a,h)anthracene	986.83		1000.55	99	
Benzo(g,h,i)perylene	933.19		1000.30	93	
Surrogate Recoveries (%)					
Naphthalene-d8	120				
Acenaphthene-d10	112				
Phenanthrene-d10	108				
Benzo(a)pyrene-d12	87				

The Business of Innovation

Client ID	GN62: North Slope Crude				
Chern ID	Ciude				
Battelle ID	BJ959NSC-P				
Sample Type	NSC				
Collection Date	01/30/07				
Extraction Date	01/30/07				
Analysis Date	02/07/07				
Analytical Instrument	MS				
% Moisture	NA				
% Lipid	NA				
Matrix	OIL				
Sample Size	5.01				
Size Unit-Basis	MG_OIL				
Minimum reporting Limit	1.3				
Units	MG/KG_OIL		Target % [Difference	Qualifier
	0007 44				
C3-Alkylbenzenes	2027.44				
C4-Alkylbenzenes	1510.05				
C5-Alkylbenzenes	795.79				
C6-Alkylbenzenes	488.99 13.52				
Benzo(b)thiophene C1-benzo(b)thiophenes	45				
	79.93		95.74	16.5	
C2-benzo(b)thiophenes C3-benzo(b)thiophenes	141.48		132.67	6.6	
C4-benzo(b)thiophenes	96.21		96.72	0.0	
Naphthalene	806.17		740.29	8.9	
C1-Naphthalenes	1634.24		1516.04	7.8	
C2-Naphthalenes	2010.9		2000.10	0.5	
C3-Naphthalenes	1445.89		1526.96	5.3	
C4-Naphthalenes	785.13		898.03	12.6	
C1-Biphenyls + Dibenzofuran	371.59		000100		
Biphenyl	240.71		220.82	9.0	
C2-Biphenyls + C1-Dibenzofurans	514.5				
Acenaphthylene		U			
Acenaphthene	13.75		14.50	5.2	
Dibenzofuran	77.57		77.75	0.2	
Fluorene	96.04		92.51	3.8	
C1-Fluorenes	223.14		227.01	1.7	
C2-Fluorenes	345.32		367.09	5.9	
C3-Fluorenes	295.4		326.32	9.5	
C4-Fluorenes	212.42				
Anthracene		U			
Phenanthrene	285.03		249.49	14.2	
C1-Phenanthrenes/Anthracenes	584.34		549.17	6.4	
C2-Phenanthrenes/Anthracenes	669.09		642.72	4.1	
C3-Phenanthrenes/Anthracenes C4-Phenanthrenes/Anthracenes	453.02		446.11	1.5	
	176.09		180.02	2.2	
Retene	74.51		210.25	44.0	
Dibenzothiophene	241.56 440.7		210.35 409.03	14.8 7.7	
C1-Dibenzothiophenes C2-Dibenzothiophenes	573.69		409.03 551.46	4.0	
C3-Dibenzothiophenes	483.6		471.36	2.6	
C4-Dibenzothiophenes	261.9		243.11	7.7	
Fluoranthene	201.0	U	240.11	1.1	
Pyrene	14.53	0	12.99	11.9	
C1-Fluoranthenes/Pyrenes	84.98		70.92	19.8	
C2-Fluoranthenes/Pyrenes	145.97		117.89	23.8	
C3-Fluoranthenes/Pyrenes	154.91		137.25	12.9	
C4-Fluoranthenes/Pyrenes	123.35				
C0-Benzo(b)naphthothiophenes	46.42				
C1-Benzo(b)naphthothiophenes	162.78				
C2-Benzo(b)naphthothiophenes	204.91				
C3-Benzo(b)naphthothiophenes	163.24				
C4-Benzo(b)naphthothiophenes	66.74				
Benzo(a)anthracene		U			

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

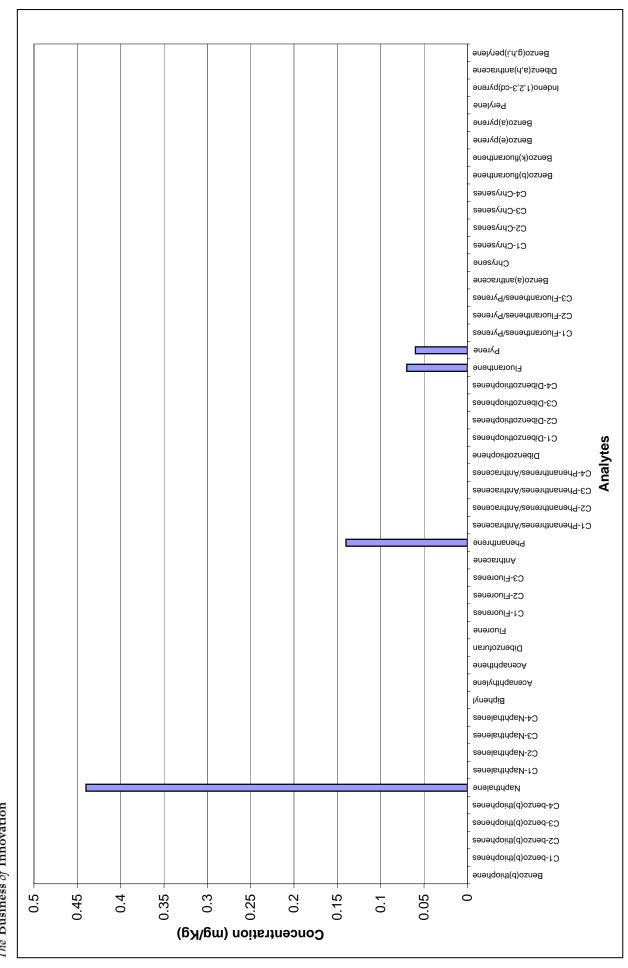
	GN62: North Slope				
Client ID	Crude				
Battelle ID	BJ959NSC-P				
Sample Type	NSC				
Collection Date	01/30/07				
Extraction Date	01/30/07				
Analysis Date	02/07/07				
Analytical Instrument	MS				
% Moisture	NA				
% Lipid	NA				
Matrix	OIL				
Sample Size	5.01				
Size Unit-Basis	MG_OIL				
Minimum reporting Limit	1.3				
Units	MG/KG_OIL		Target % D	oifference	Qualifier
Chrysene	51.12		47.18	8.4	
C1-Chrysenes	85.28		78.82	8.2	
C2-Chrysenes	113.52		102.67	10.6	
C3-Chrysenes	97.23		85.36	13.9	
C4-Chrysenes	62.06		61.99	0.1	
Benzo(b)fluoranthene	6.18		6.08	1.6	
Benzo(k)fluoranthene		U			
Benzo(e)pyrene	13.86		12.88	7.6	
Benzo(a)pyrene		U			
Perylene		U			
		U			
Indeno(1,2,3-cd)pyrene		J			
	1.1	0			
Indeno(1,2,3-cd)pyrene Dibenz(a,h)anthracene Benzo(g,h,i)perylene	1.1 3.86	U	3.44	12.2	

Surrogate Recoveries (%)

Naphthalene-d8	109
Acenaphthene-d10	106
Phenanthrene-d10	97
Benzo(a)pyrene-d12	119

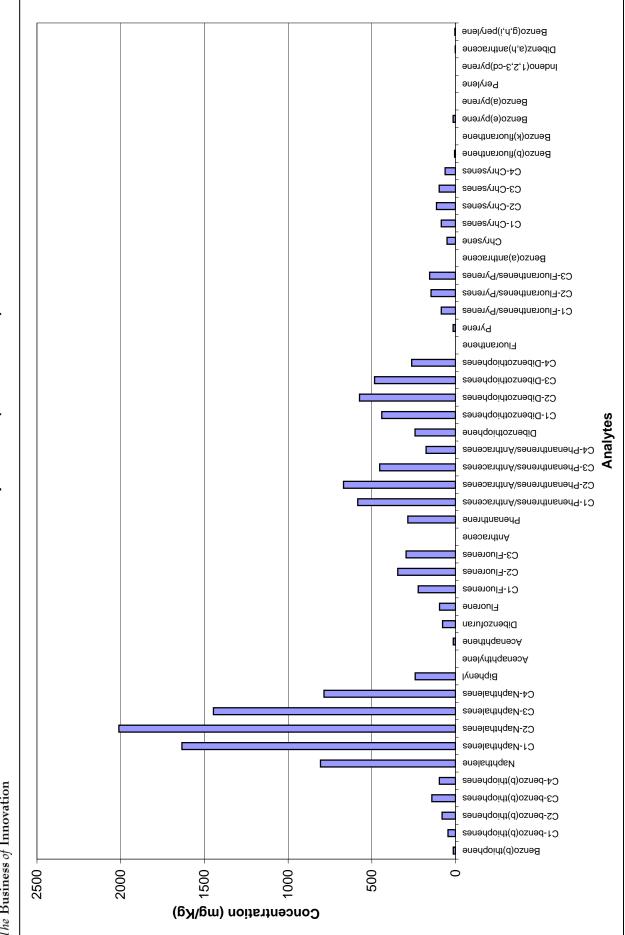
Baltelle The Business of Innovation

Procedural Blank (BJ941PB-P)



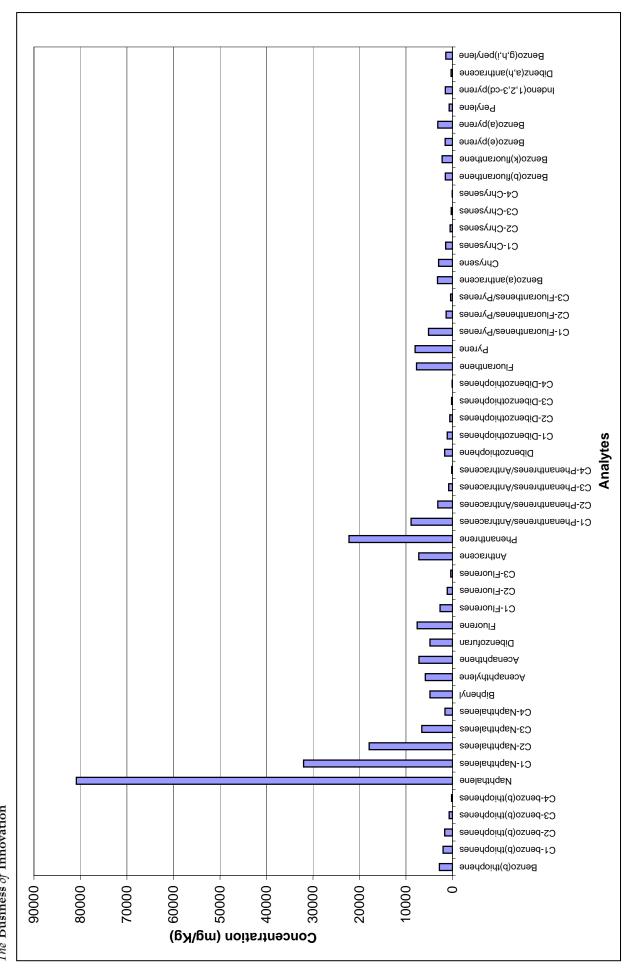


GN62: North Slope Crude (BJ959NSC-P)



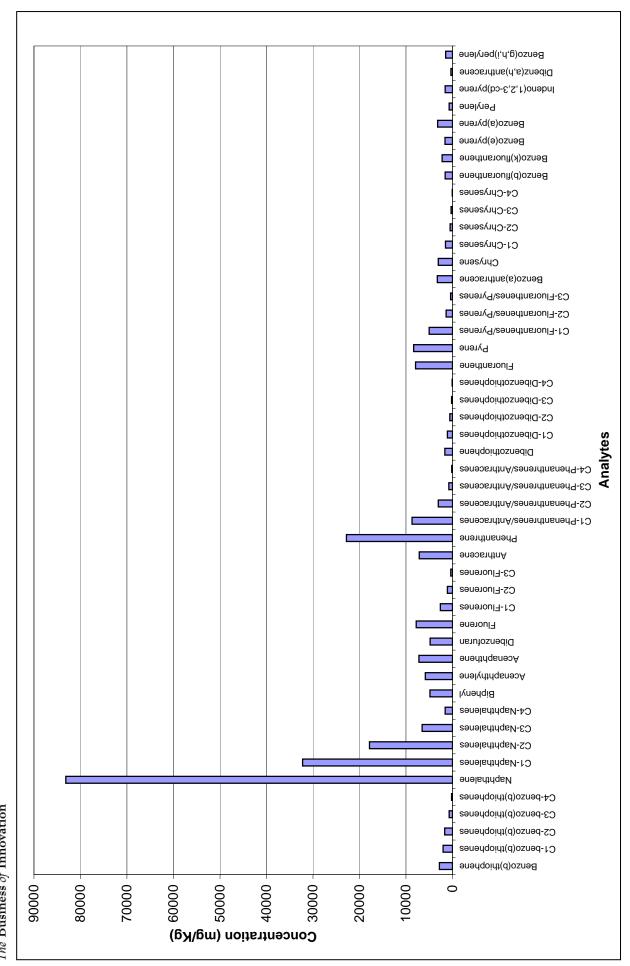
Baffelle The Business of Innovation

GWP07DW401 (R5522-P)



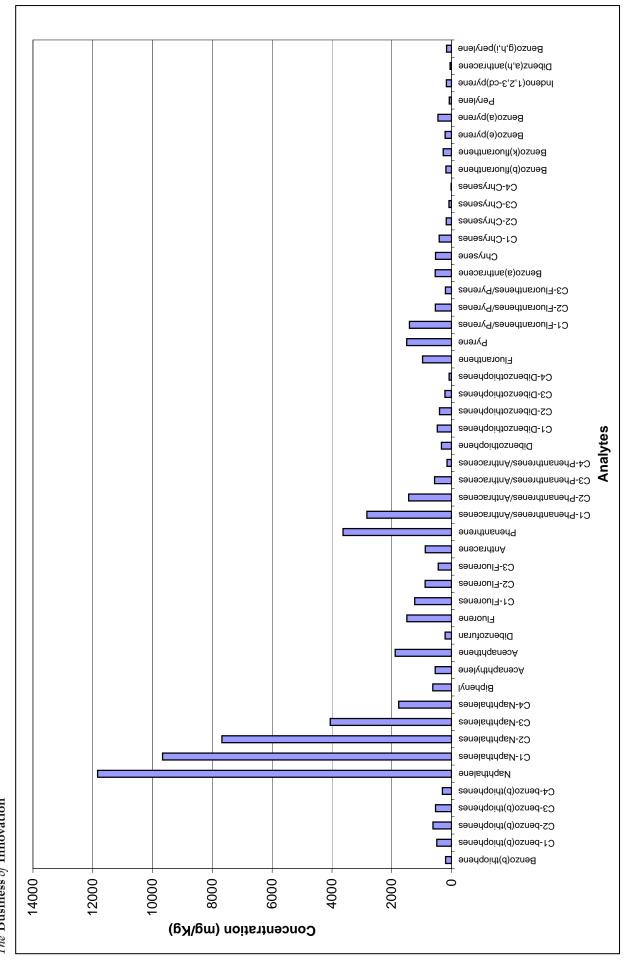
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GWP07DW402 (R5523-P)



Baltelle The Business of Innovation

GWP07MW9 (R5524-P)



The Business of Innovation

Client ID	GWP07DW401	GWP07DW402	GWP07MW9
Battelle ID	R5522-P	R5523-P	R5524-P
Sample Type	SA	SA	SA
Collection Date	01/11/07	01/11/07	01/12/07
Extraction Date	01/30/07	01/30/07	01/30/07
Analysis Date	02/09/07	02/09/07	02/09/07
Analytical Instrument	02/09/07 MS	02/09/07 MS	02/09/07 MS
5	-		-
% Moisture	NA NA	NA NA	NA NA
% Lipid			
Matrix	NAPL 40.50	NAPL 47.40	NAPL 45.80
Sample Size	49.50		45.80
Size Unit-Basis	MG_OIL	MG_OIL	MG_OIL
Minimum Reporting Limit	1.39	1.45	1.36
Units	MG/KG_OIL	MG/KG_OIL	MG/KG_OIL
C23 Tricyclic Terpane	U	U	40.62
C29 Tricyclic Terpane -22S	Ŭ	Ŭ	12.03
C29 Tricyclic Terpane -22R	Ŭ	Ŭ	11.13
18a(H)-22,29,30-Trisnorneohopane -TS	Ŭ	Ŭ	7.86
17a(H)-22,29,30-Trisnorhopane -TM	Ŭ	Ŭ	10.84
30-Norhopane	6.51	6.04	29.84
18a(H) & 18b(H)-Oleananes	U	U	12.11
Hopane	10.38	11.12	57.14
30-Homohopane -22S	U	U	13.98
30-Homohopane -22R	U	U	10.97
13b(H),17a(H)-20S-Diacholestane	U	U	65.16
13b(H),17a(H)-20R-Diacholestane	U	U	40.91
14a(H),17a(H)-20R-methylcholestane	6.86	7.91	86.78
14a(H),17a(H)-20S-Ethylcholestane	3.58	3.08	29.66
14a(H),17a(H)-20R-Ethylcholestane	3.39	4.74	47.85
C21-TAS	U	U	9.32
C26-TAS(20S)	1.44	1.6	14.27
C26,C27-TAS	4.78	5	46.72
C27-TAS(20R)	2.36	2.57	25.18
C28-TAS(20S)	1.6	1.68	13.93
C28-TAS(20R)	1.38 J	1.71	11.88
C21-MAS	U	U	U
C22-MAS	U	U	U
C27-MAS	U	U	U
C27-20R-MAS	U	U	10.16
C27-20S-MAS	U	U	9.31
C28-20S-MAS	U	U	21.72
C27-C2920S/R-MAS	U	U	27.59
C29-20S-MAS	U	U	7.49
C29-20R-MAS	U	U	17.37
TAS_245	NA	NA	NA
MAS_239	NA	NA	NA
Surrogate Recoveries (%)			

5b(H)-Cholane	570 NME	568 NME	112

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID Procedural E	lank
Battelle ID BJ941	PB-P
Sample Type	PB
Collection Date 01/3	0/07
Extraction Date 01/3	0/07
Analysis Date 02/1	2/07
Analytical Instrument	MS
% Moisture	NA
% Lipid	NA
Matrix	APL
Sample Size	5.00
Size Unit-Basis MG	_OIL
Minimum Reporting Limit	1.31
Units MG/KG	_OIL

C23 Tricyclic Terpane	U
C29 Tricyclic Terpane -22S	U
C29 Tricyclic Terpane -22R	U
18a(H)-22,29,30-Trisnorneohopane -TS	U
17a(H)-22,29,30-Trisnorhopane -TM	U
30-Norhopane	U
18a(H) & 18b(H)-Oleananes	U
Hopane	U
30-Homohopane -22S	U
30-Homohopane -22R	U
13b(H),17a(H)-20S-Diacholestane	U
13b(H),17a(H)-20R-Diacholestane	U
14a(H),17a(H)-20R-methylcholestane	U
14a(H),17a(H)-20S-Ethylcholestane	U
14a(H),17a(H)-20R-Ethylcholestane	U
C21-TAS	U
C26-TAS(20S)	U
C26,C27-TAS	U
C27-TAS(20R)	U
C28-TAS(20S)	U
C28-TAS(20R)	U
C21-MAS	U
C22-MAS	U
C27-MAS	U
C27-20R-MAS	U
C27-20S-MAS	U
C28-20S-MAS	U
C27-C2920S/R-MAS	U
C29-20S-MAS	U
C29-20R-MAS	U
TAS_245	NA
MAS_239	NA

Surrogate Recoveries (%)

5b(H)-Cholane

The Business of Innovation

Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

Client ID	Laboratory Control Sample	
Client ID	Sample	
Battelle ID	BJ942LCS-P	
Sample Type	LCS	
Collection Date	01/30/07	
Extraction Date	01/30/07	
Analysis Date	02/12/07	
Analytical Instrument	MS	
% Moisture	NA	
% Lipid	NA	
Matrix	NAPL	
Sample Size	NA	
Size Unit-Basis	NA	
Minimum Reporting Limit	7.03	
Units	NG	Target % Recovery Qualifier
		ů ,
C23 Tricyclic Terpane	U	
C29 Tricyclic Terpane -22S	U	
C29 Tricyclic Terpane -22R	U	
18a(H)-22,29,30-Trisnorneohopane -TS	U	
17a(H)-22,29,30-Trisnorhopane -TM	U	
30-Norhopane	U	
18a(H) & 18b(H)-Oleananes	U	
Hopane	U	
30-Homohopane -22S	U	
30-Homohopane -22R	U	
13b(H),17a(H)-20S-Diacholestane	U	
13b(H),17a(H)-20R-Diacholestane	U	
14a(H),17a(H)-20R-methylcholestane	U	
14a(H),17a(H)-20S-Ethylcholestane	U	
14a(H),17a(H)-20R-Ethylcholestane	U	
C21-TAS	U	
C26-TAS(20S)	U	
C26,C27-TAS	U	
C27-TAS(20R)	U	
C28-TAS(20S)	U	
C28-TAS(20R)	U	
C21-MAS	U	
C22-MAS	U	
C27-MAS	U	
C27-20R-MAS	U	
C27-20S-MAS	U	
C28-20S-MAS	U	
C27-C2920S/R-MAS	U	
C29-20S-MAS	U	
C29-20R-MAS	U	
TAS_245	NA	
MAS_239	NA	

Surrogate Recoveries (%)

5b(H)-Cholane

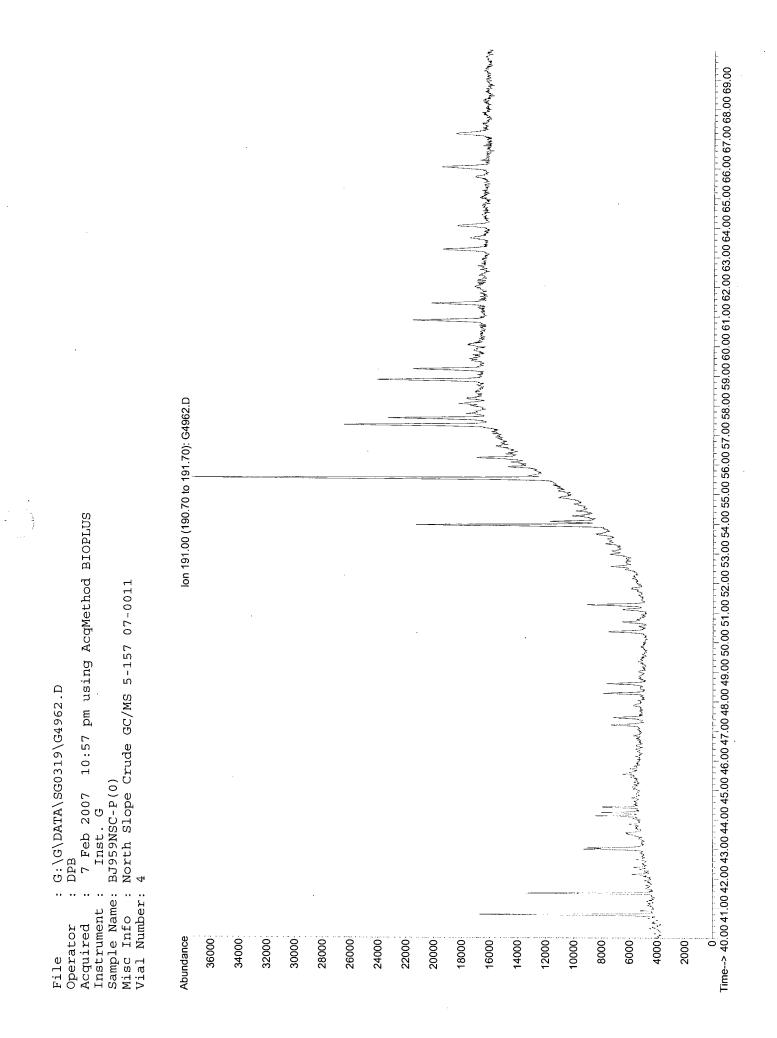
The Business of Innovation

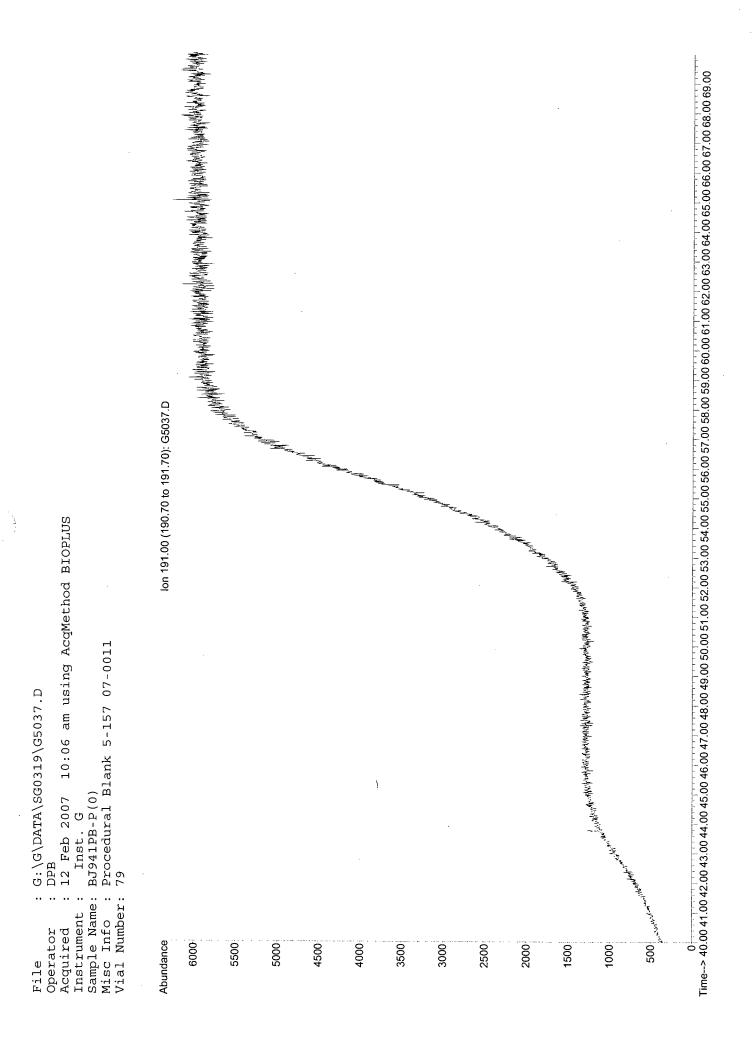
Project Client: Exponent, Inc. Project Name: Exponent - Gas Works Park Project Number: N106746-0001

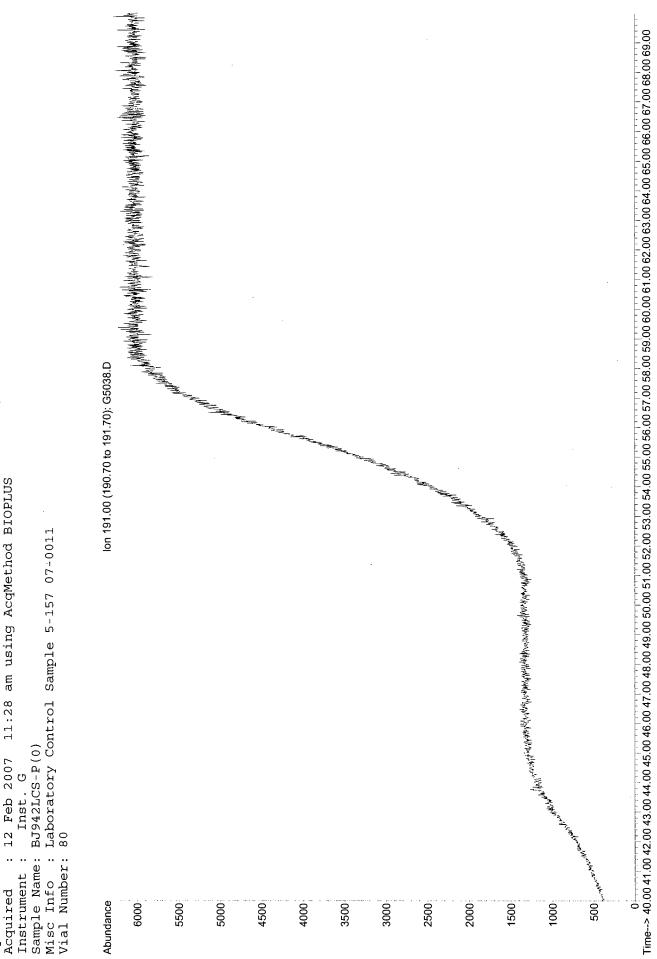
Client ID Crude Battelle ID BJ959NSC-P Sample Type NSC Collection Date 01/30/07 Extraction Date 02/07/07 Analytical Instrument MS % Moisture NA Matrix OlL Sample Size 5.01 Size Unit-Basis MG/G_OLL Minimum Reporting Limit 1.3 Units MG/KG_OLL Target % Difference C26-TAS(20S) 15.53 C26,C27-TAS 55.26 C27-TAS(20R) 31.26 C28-TAS(20R) 31.26 C28-TAS(20R) 31.26 C27-AS 55.26 C27-TAS 55.26 C27-TAS 55.26 C27-TAS 55.26 C27-TAS 55.26 C27-MAS		GN62: North Slope	
Sample Type NSC Collection Date 01/30/07 Extraction Date 01/30/07 Analysis Date 02/07/07 Analysis Date NA % Moisture NA Matrix OIL Sample Size 5.01 Size Unit-Basis MG_OIL Minimum Reporting Limit 1.3 Units MG/KG_OIL Target % Difference Qualifier C21-TAS 15.53 C26, C27-TAS 55.26 C27-TAS(20S) 31.48 C28-TAS(20R) 31.48 C28-TAS(20R) 31.48 C28-TAS(20R) 31.48 C27-C29LMAS 3.54 C27-C29LMAS 2.57 C27-AS 5.98 2.57 C27-C29LMAS C27-C29LOS-MAS	Client ID	Crude	
Sample Type NSC Collection Date 01/30/07 Extraction Date 01/30/07 Analysis Date 02/07/07 Analysis Date NA % Moisture NA Matrix OIL Sample Size 5.01 Size Unit-Basis MG_OIL Minimum Reporting Limit 1.3 Units MG/KG_OIL Target % Difference Qualifier C21-TAS 15.53 C26, C27-TAS 55.26 C27-TAS(20S) 31.48 C28-TAS(20R) 31.48 C28-TAS(20R) 31.48 C28-TAS(20R) 31.48 C27-C29LMAS 3.54 C27-C29LMAS 2.57 C27-AS 5.98 2.57 C27-C29LMAS C27-C29LOS-MAS			
Collection Date 01/30/07 Analysis Date 02/07/07 Analysical Instrument MS % Moisture NA Matrix OIL Sample Size 5.01 Size Unit-Basis MG_OIL Minimum Reporting Limit 1.3 Units MG/KG_OIL Target % Difference C26-TAS(20S) 15.53 C26-C27-TAS 55.26 C27-TAS(20R) 37.91 C28-TAS(20S) 31.48 C27-TAS 55.26 C27-MAS 3.54 C27-MAS 4.73 C27-20S-MAS 5.98 C27-20S-MAS 14.9 C28-Z0S-MAS 14.9 C27-C2920S/R-MAS 12.33 C29-20R-MAS 9.14 TAS_245 NA			
Extraction Date 01/30/07 Analytical Instrument MS % Moisture NA % Lipid NA Matrix OlL Sample Size 5.01 Size Unit-Basis MG_OIL Minimum Reporting Limit 1.3 Units MG/KG_OIL Target % Difference Qualifier C21-TAS 18.59 C26-TAS(20S) 15.53 C26,C27-TAS 55.26 C27-TAS(20R) 37.91 C28-TAS(20R) 31.48 C27-MAS 6.09 C27-MAS 3.54 C27-MAS 5.98 C27-C27-QS-MAS 2.57 C28-DS-MAS 14.9 C27-C2920S/R-MAS 12.33 C29-20R-MAS 9.14 TAS_245 NA			
Analysis Date 02/07/07 Analytical Instrument MS % Moisture NA % Lipid NA Matrix OIL Sample Size 5.01 Size Unit-Basis MG_OIL Minimum Reporting Limit 1.3 Units MG/KG_OIL Target % Difference C21-TAS 18.59 C26-TAS(20S) 15.53 C26,C27-TAS 55.26 C27-TAS(20R) 37.91 C28-TAS(20R) 31.48 C28-TAS(20R) 31.48 C28-TAS(20R) 31.48 C22-MAS 5.98 C27-MAS 4.73 C27-OR-MAS 5.98 C27-C2920S/R-MAS 14.9 C27-C2920S/R-MAS 4.08 C29-20R-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA			
Analytical Instrument MS % Moisture NA % Lipid NA % Lipid NA Matrix OIL Sample Size 5.01 Size Unit-Basis MG_OIL Minimum Reporting Limit 1.3 Units MG/KG_OIL Target % Difference C21-TAS 15.53 C26-TAS(20S) 15.53 C26-TAS(20S) 15.53 C26-TAS(20R) 37.91 C28-TAS(20R) 31.48 C22-MAS 6.09 C22-MAS 3.54 C27-MAS 4.73 C27-20R-MAS 5.98 C27-20S-MAS 14.9 C28-C29S/R-MAS 14.9 C29-20S-MAS 4.08 C29-20S-MAS 9.14 TAS_245 NA	Extraction Date	01/30/07	
% Moisture NA % Lipid NA Matrix OIL Sample Size 5.01 Size Unit-Basis MG_OIL Minimum Reporting Limit 1.3 Units MG/KG_OIL Target % Difference Qualifier C21-TAS 18.59 C26-TAS(20S) 15.53 C26-C27-TAS 55.26 C27-TAS(20R) 37.91 C28-TAS(20S) 31.48 C28-TAS(20R) 31.26 C21-MAS 6.09 C22-MAS 3.54 C27-TAS 5.98 C27-TAS 2.57 C28-DS-MAS 14.9 C27-C2920S/R-MAS 12.33 C29-20R-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	Analysis Date	02/07/07	
% Lipid NA Matrix OIL Sample Size 5.01 Size Unit-Basis MG_OIL Minimum Reporting Limit 1.3 Units MG/KG_OIL Target % Difference Qualifier C21-TAS 18.59 C26-TAS(20S) 15.53 C26,C27-TAS 55.26 C27-TAS(20S) 31.48 C28-TAS(20R) 31.26 C21-MAS 6.09 C22-MAS 3.54 C27-CAS 5.98 C27-C20S-MAS 2.57 C28-SOS 14.9 C27-C2920S/R-MAS 12.33 C29-20R-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	Analytical Instrument	MS	
Matrix OIL Sample Size 5.01 Size Unit-Basis MG_OIL Minimum Reporting Limit 1.3 Units MG/KG_OIL Target % Difference C21-TAS 18.59 C26-TAS(20S) 15.53 C26-TAS(20R) 37.91 C28-TAS(20R) 31.48 C28-TAS(20R) 31.48 C21-MAS 6.09 C22-MAS 3.54 C27-TAS 5.58 C27-TAS 5.54 C28-TAS(20R) 31.26 C21-MAS 6.09 C22-MAS 3.54 C27-OR-MAS 5.98 C27-C20S-MAS 14.9 C27-C2920S/R-MAS 14.9 C27-C2920S/R-MAS 14.9 C29-20R-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	% Moisture	NA	
Sample Size 5.01 Size Unit-Basis MG_OLL Minimum Reporting Limit 1.3 Units MG/KG_OLL Target % Difference Qualifier C21-TAS 18.59 C26-TAS(20S) 15.53 C26-C27-TAS 55.26 C27-TAS(20R) 37.91 C28-TAS(20S) 31.48 C28-TAS(20R) 31.26 C21-MAS 6.09 C27-MAS 3.54 C27-OR-MAS 4.73 C27-20S-MAS 2.57 C28-20S-MAS 14.9 C27-C2920S/R-MAS 12.33 C29-20R-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	% Lipid	NA	
Size Unit-Basis MG_OIL Minimum Reporting Limit 1.3 Units MG/KG_OIL Target % Difference Qualifier C21-TAS 18.59 C26-TAS(20S) 15.53 C26,C27-TAS 55.26 C27-TAS(20R) 37.91 C28-TAS(20S) 31.48 C28-TAS(20R) 31.26 C21-MAS 6.09 C27-VAS 3.54 C27-VAS 4.73 C27-20R-MAS 5.98 C27-20S-MAS 14.9 C27-C2920S/R-MAS 14.9 C27-02S-MAS 4.08 C29-20S-MAS 4.08 C29-20S-MAS 9.14 TAS_245 NA	Matrix	OIL	
Minimum Reporting Limit 1.3 MG/KG_OIL Target % Difference Qualifier C21-TAS 18.59 </td <td>Sample Size</td> <td>5.01</td> <td></td>	Sample Size	5.01	
Units MG/KG_OIL Target % Difference Qualifier C21-TAS 18.59	Size Unit-Basis	MG_OIL	
C21-TAS 18.59 C26-TAS(20S) 15.53 C26,C27-TAS 55.26 C27-TAS(20R) 37.91 C28-TAS(20S) 31.48 C28-TAS(20R) 31.26 C21-MAS 6.09 C22-MAS 3.54 C27-WAS 4.73 C27-20R-MAS 5.98 C27-20S-MAS 2.57 C28-20S-MAS 14.9 C27-C2920S/R-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	Minimum Reporting Limit	1.3	
C26-TAS(20S) 15.53 C26,C27-TAS 55.26 C27-TAS(20R) 37.91 C28-TAS(20S) 31.48 C28-TAS(20R) 31.26 C21-MAS 6.09 C22-MAS 3.54 C27-QR-MAS 5.98 C27-20S-MAS 2.57 C28-QS-MAS 14.9 C29-20S/R-MAS 12.33 C29-20R-MAS 9.14 TAS_245 NA	Units	MG/KG_OIL	Target % Difference Qualifier
C26-TAS(20S) 15.53 C26,C27-TAS 55.26 C27-TAS(20R) 37.91 C28-TAS(20S) 31.48 C28-TAS(20R) 31.26 C21-MAS 6.09 C22-MAS 3.54 C27-QR-MAS 5.98 C27-20S-MAS 2.57 C28-QS-MAS 14.9 C29-20S/R-MAS 12.33 C29-20R-MAS 9.14 TAS_245 NA			
C26,C27-TAS 55.26 C27-TAS(20R) 37.91 C28-TAS(20S) 31.48 C28-TAS(20R) 31.26 C21-MAS 6.09 C22-MAS 3.54 C27-OR-MAS 5.98 C27-20S-MAS 2.57 C28-C2920S/R-MAS 14.9 C29-20S-MAS 12.33 C29-20R-MAS 9.14 TAS_245 NA	C21-TAS	18.59	
C27-TAS(20R) 37.91 C28-TAS(20S) 31.48 C28-TAS(20R) 31.26 C21-MAS 6.09 C22-MAS 3.54 C27-VAS 4.73 C27-20R-MAS 5.98 C27-20S-MAS 2.57 C28-20S-MAS 14.9 C29-20S-MAS 12.33 C29-20S-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	C26-TAS(20S)	15.53	
C28-TAS(20S) 31.48 C28-TAS(20R) 31.26 C21-MAS 6.09 C22-MAS 3.54 C27-MAS 4.73 C27-20R-MAS 5.98 C27-20S-MAS 2.57 C28-20S-MAS 14.9 C27-C2920S/R-MAS 4.08 C29-20S-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	C26,C27-TAS	55.26	
C28-TAS(20R) 31.26 C21-MAS 6.09 C22-MAS 3.54 C27-MAS 4.73 C27-20R-MAS 5.98 C27-20S-MAS 2.57 C28-20S-MAS 14.9 C27-C2920S/R-MAS 4.08 C29-20S-MAS 9.14 TAS_245 NA	C27-TAS(20R)	37.91	
C21-MAS 6.09 C22-MAS 3.54 C27-MAS 4.73 C27-20R-MAS 5.98 C27-20S-MAS 2.57 C28-20S-MAS 14.9 C27-C2920S/R-MAS 12.33 C29-20S-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	C28-TAS(20S)	31.48	
C22-MAS 3.54 C27-MAS 4.73 C27-20R-MAS 5.98 C27-20S-MAS 2.57 C28-20S-MAS 14.9 C27-C2920S/R-MAS 12.33 C29-20S-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	C28-TAS(20R)	31.26	
C27-MAS 4.73 C27-20R-MAS 5.98 C27-20S-MAS 2.57 C28-20S-MAS 14.9 C27-C2920S/R-MAS 12.33 C29-20S-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	C21-MAS	6.09	
C27-20R-MAS 5.98 C27-20S-MAS 2.57 C28-20S-MAS 14.9 C27-C2920S/R-MAS 12.33 C29-20S-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	C22-MAS	3.54	
C27-20S-MAS 2.57 C28-20S-MAS 14.9 C27-C2920S/R-MAS 12.33 C29-20S-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	C27-MAS	4.73	
C28-20S-MAS 14.9 C27-C2920S/R-MAS 12.33 C29-20S-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	C27-20R-MAS	5.98	
C27-C2920S/R-MAS 12.33 C29-20S-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	C27-20S-MAS	2.57	
C29-20S-MAS 4.08 C29-20R-MAS 9.14 TAS_245 NA	C28-20S-MAS	14.9	
C29-20R-MAS 9.14 TAS_245 NA	C27-C2920S/R-MAS	12.33	
TAS_245 NA	C29-20S-MAS	4.08	
	C29-20R-MAS	9.14	

Surrogate Recoveries (%)

5b(H)-Cholane







1.47

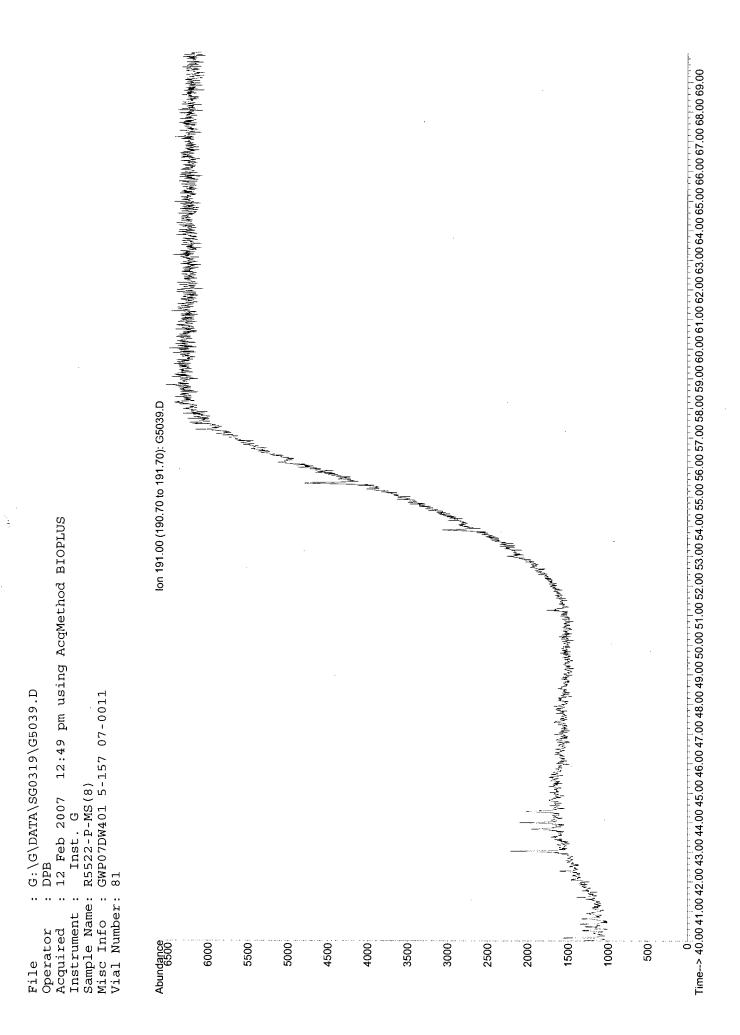
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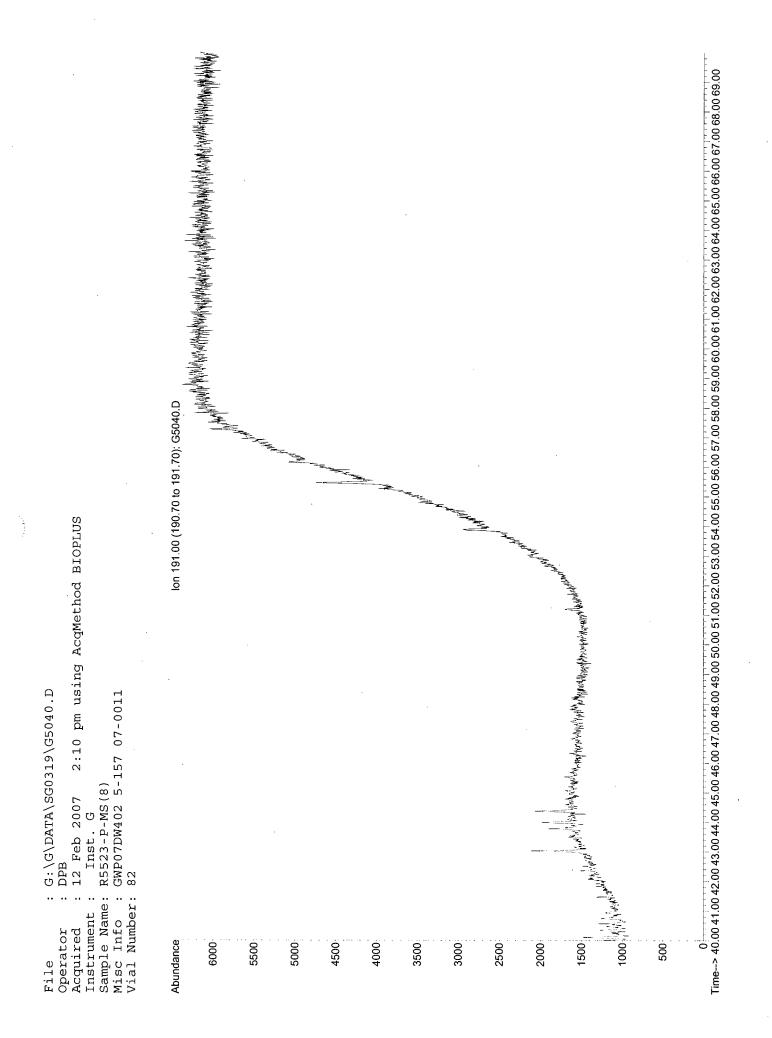
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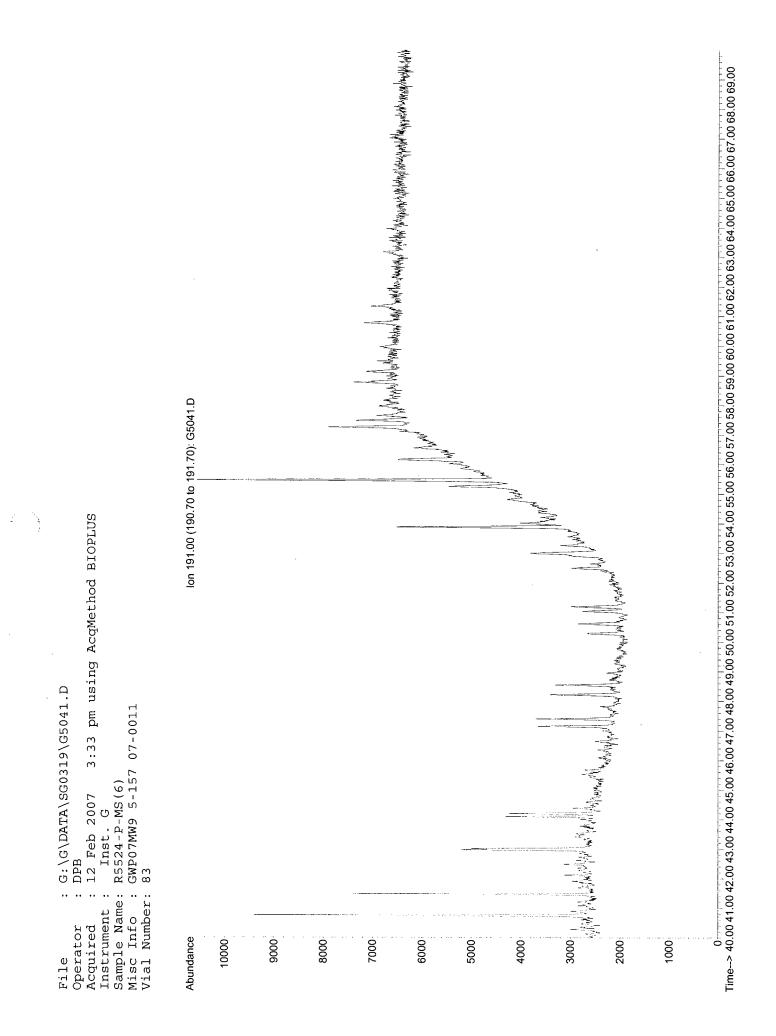
12 Feb 2007

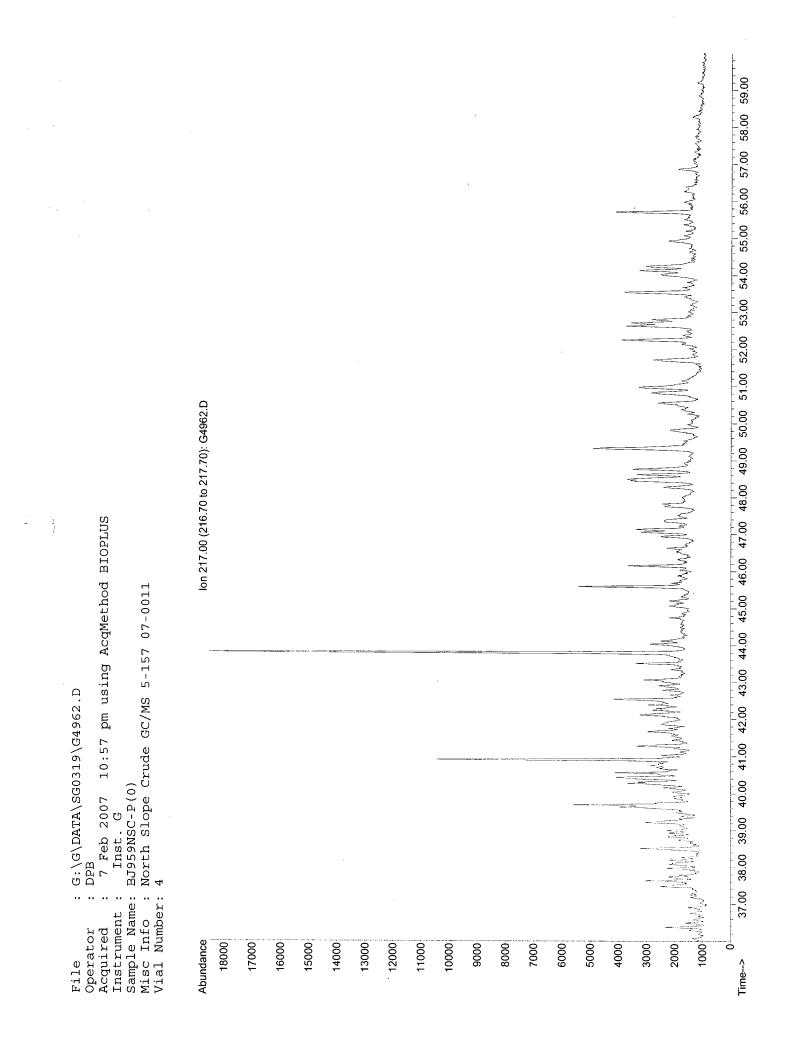
Operator

File





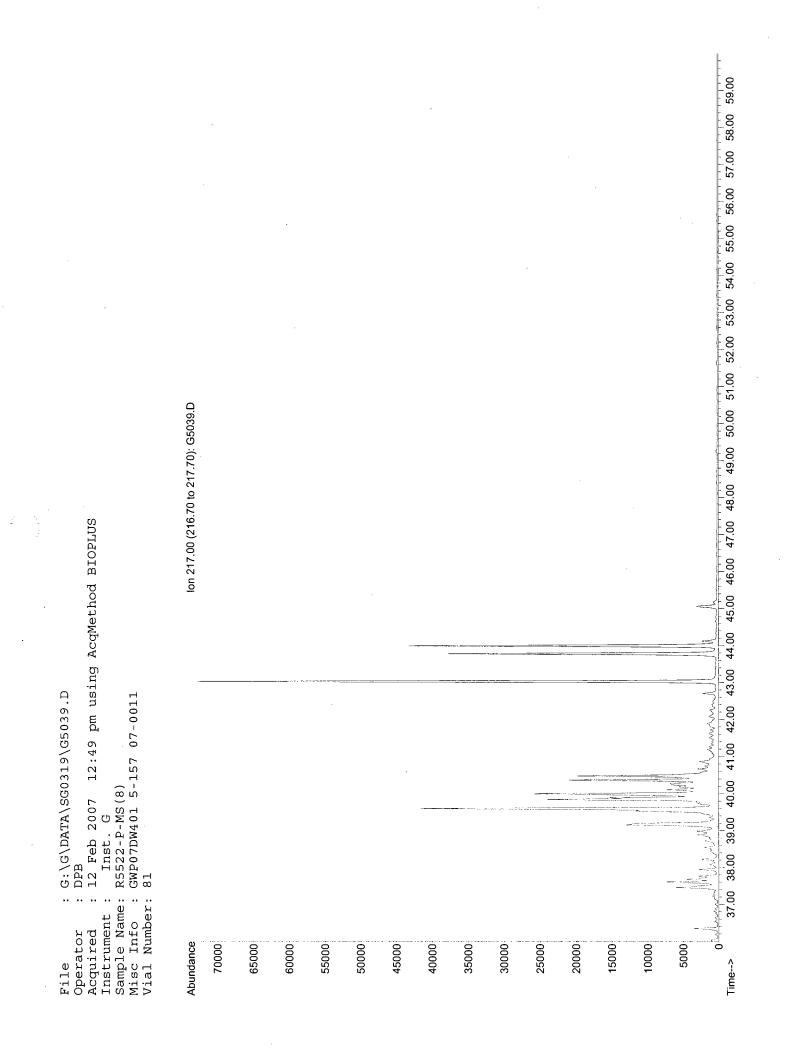


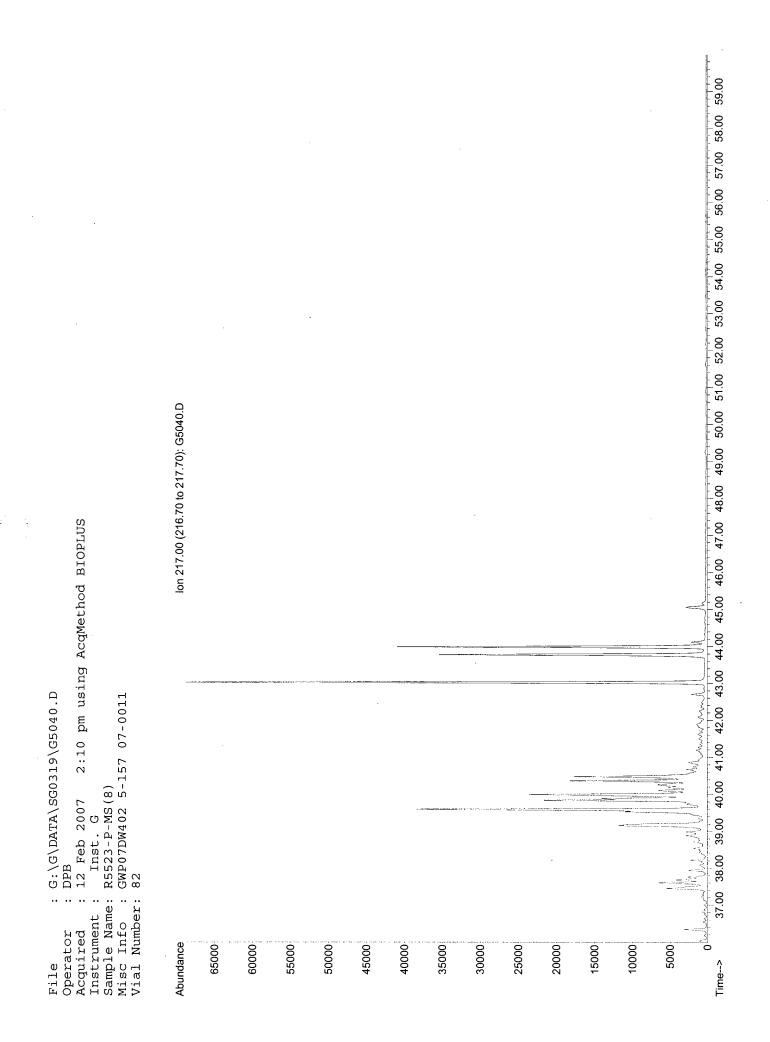


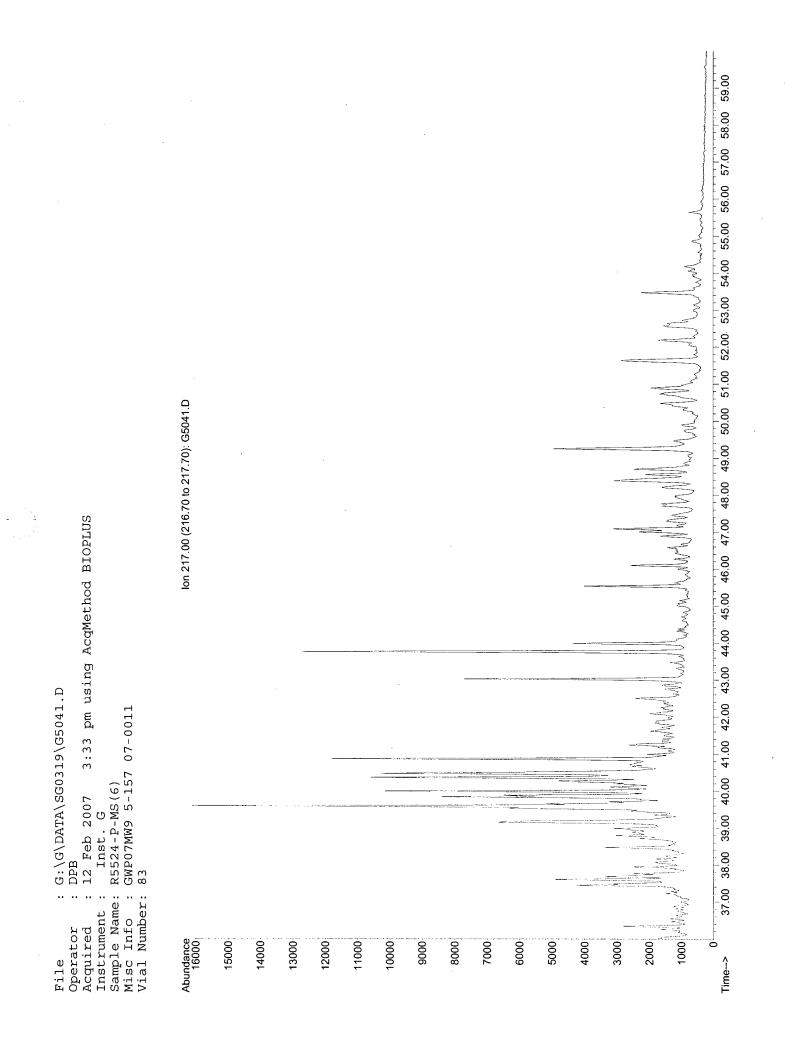
37.00 38.00 39.00 41.00 41.00 42.00 43.00 44.00 45.00 47.00 48.00 49.00 50.00 51.00 52.00 53.00 56.00 56.00 57.00 58.00 59.00 lon 217.00 (216.70 to 217.70); G5037.D 10:06 am using AcqMethod BIOPLUS File : G:\G\DATA\SG0319\G5037.D Operator : DPB Acquired : 12 Feb 2007 10:06 am using Ac Instrument : Inst. G Sample Name: BJ941PB-P(0) Misc Info : Procedural Blank 5-157 07-0011 Vial Number: 79 Abundance Time-->

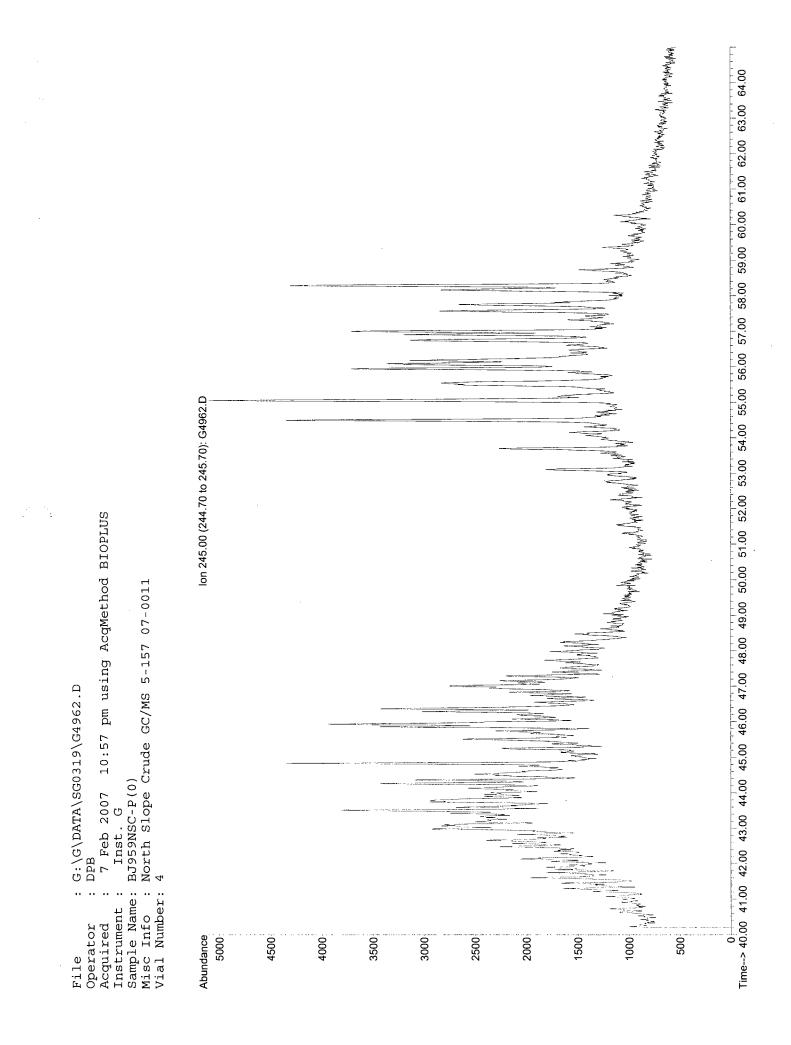
lon 217.00 (216.70 to 217.70): G5038.D File : G:\G\DATA\SG0319\G5038.D Operator : DPB Acquired : 12 Feb 2007 11:28 am using AcqMethod BIOPLUS Instrument : Inst. G Sample Name: BJ942LCS-P(0) Misc Info : Laboratory Control Sample 5-157 07-0011 Vial Number: 80 Abundance

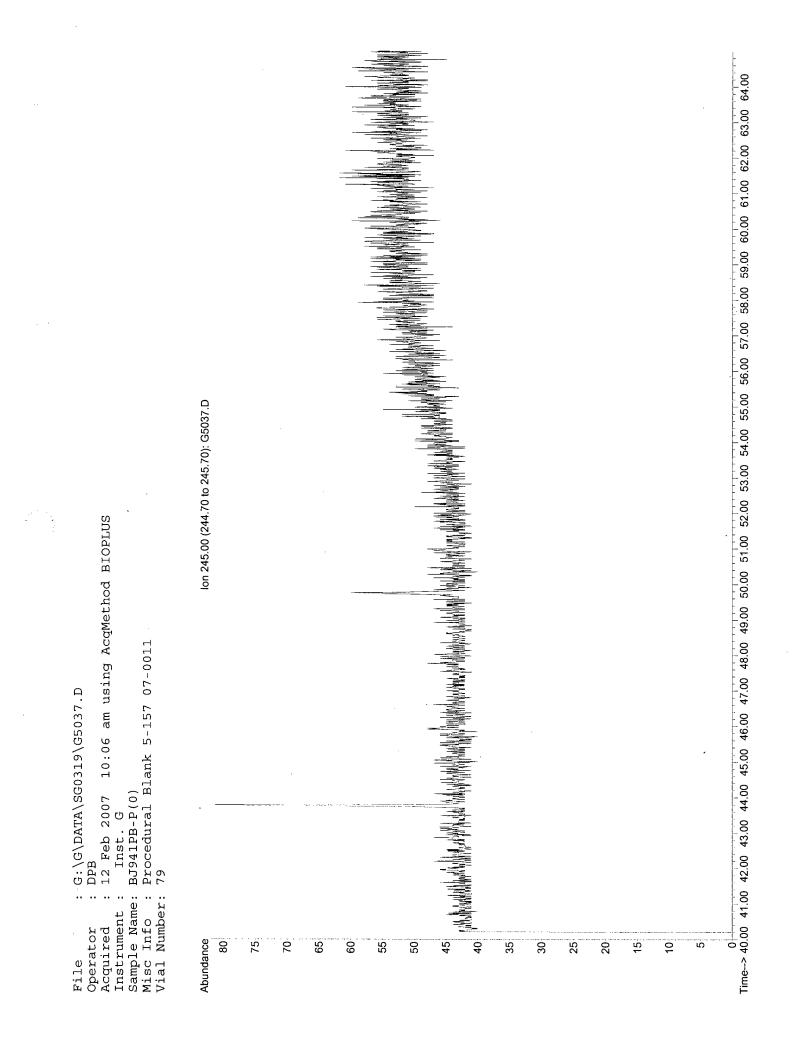
37.00 38.00 39.00 40.00 41.00 42.00 43.00 44.00 45.00 47.00 48.00 49.00 50.00 51.00 52.00 53.00 55.00 56.00 57.00 58.00 59.00 Time-->

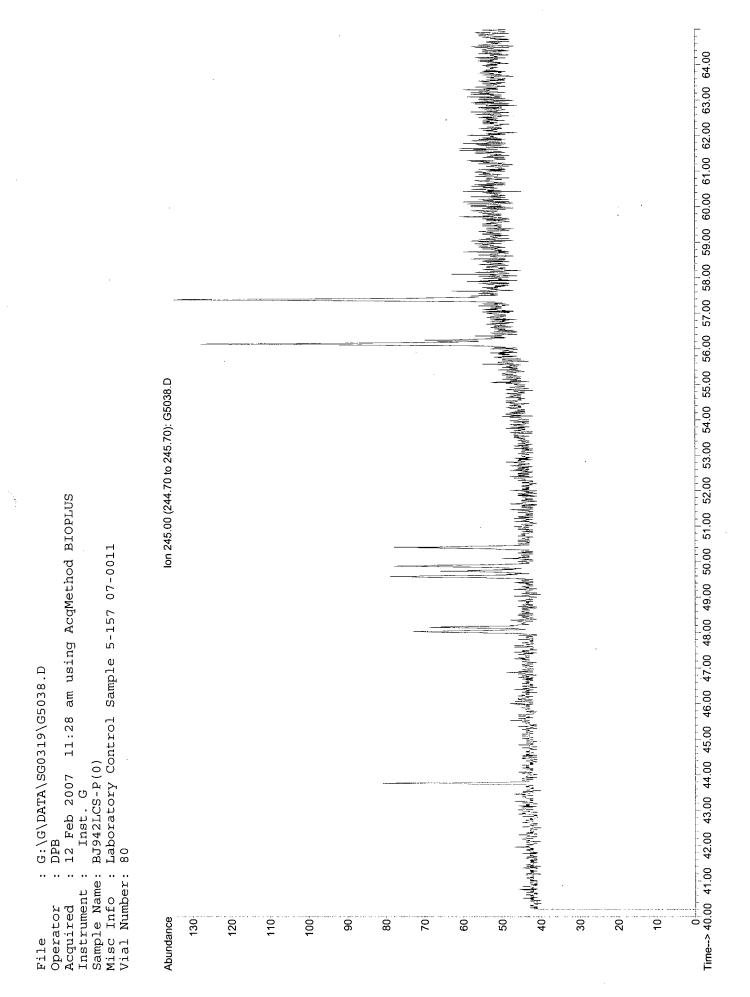


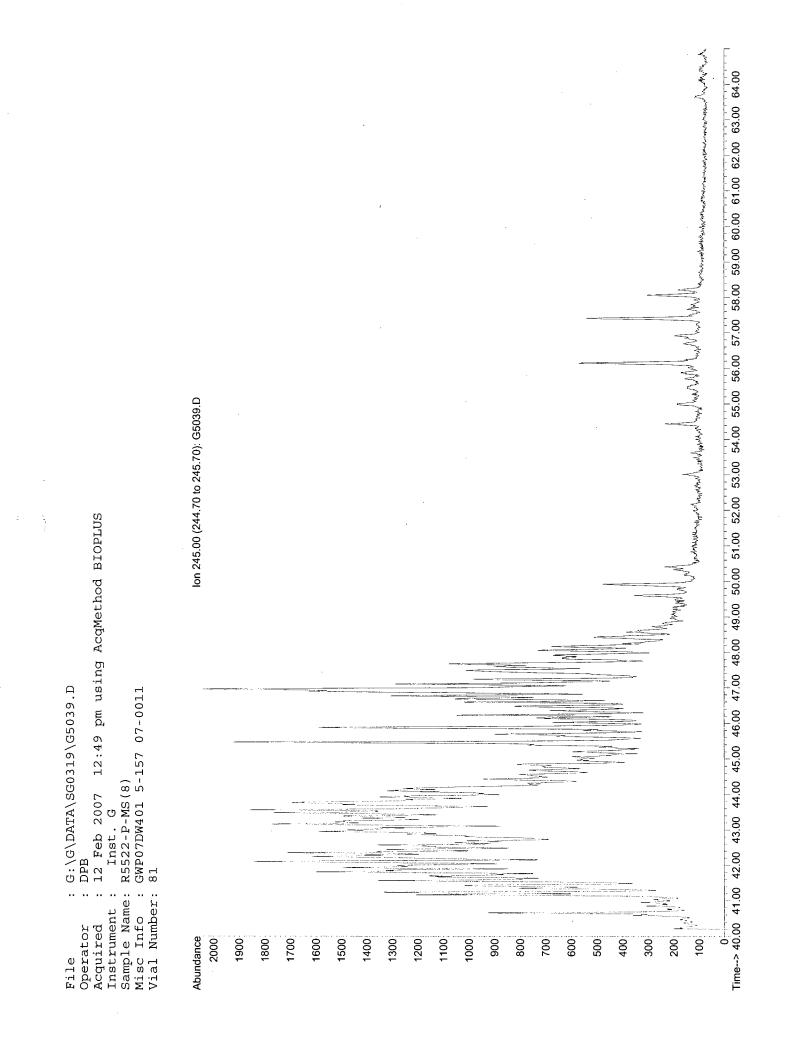


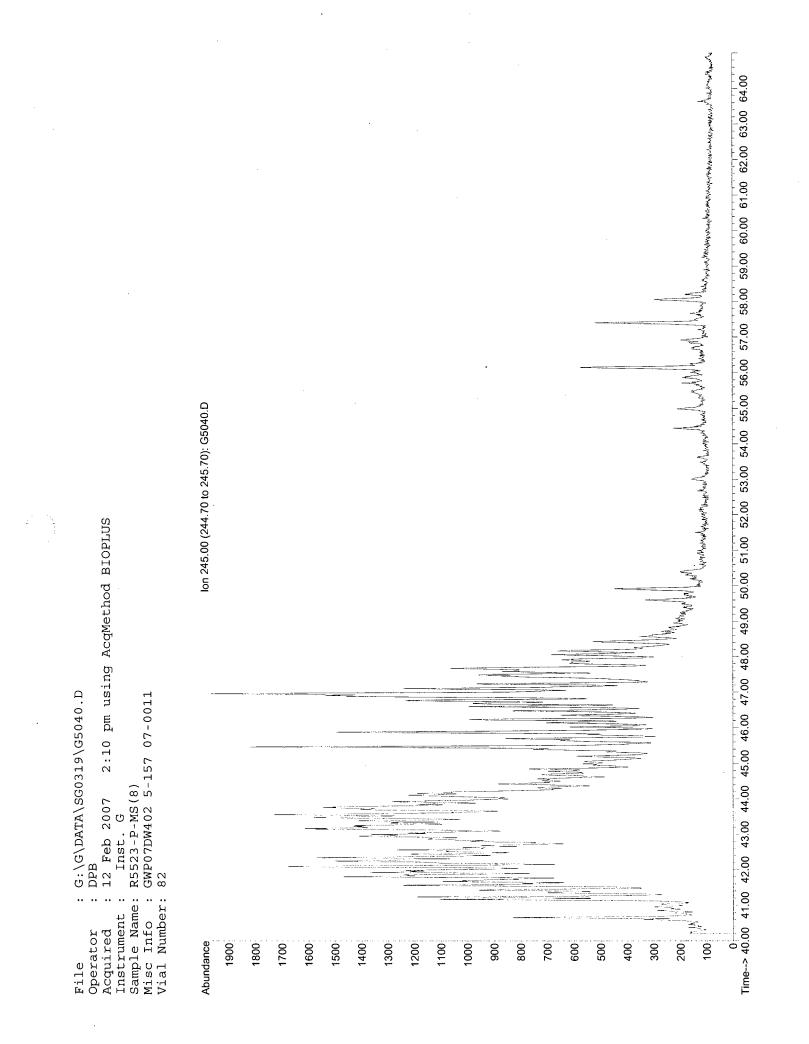


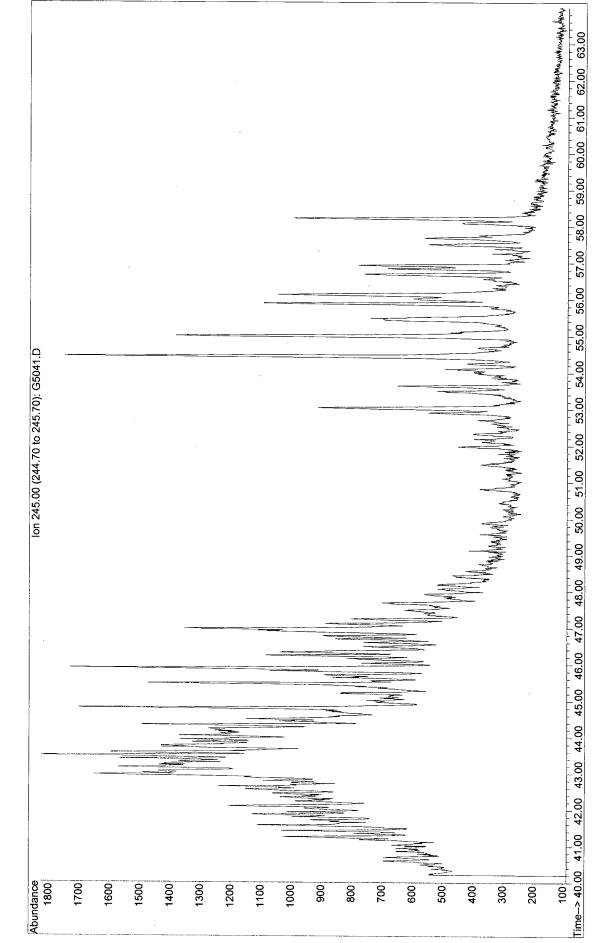




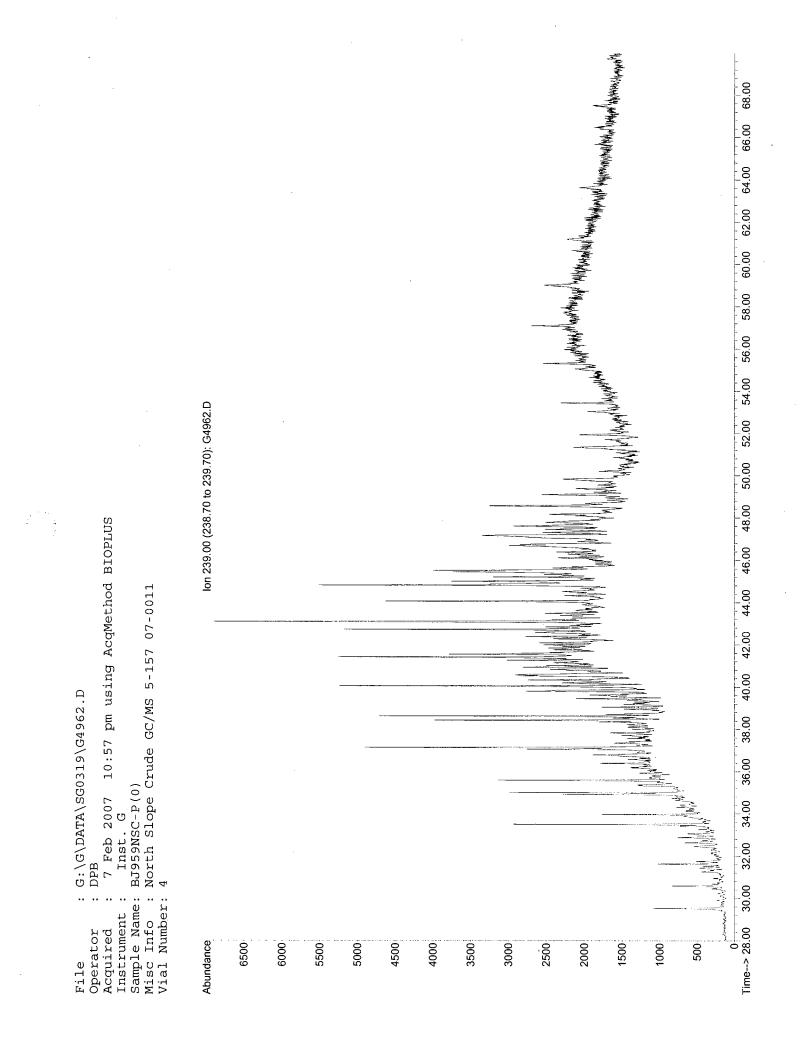


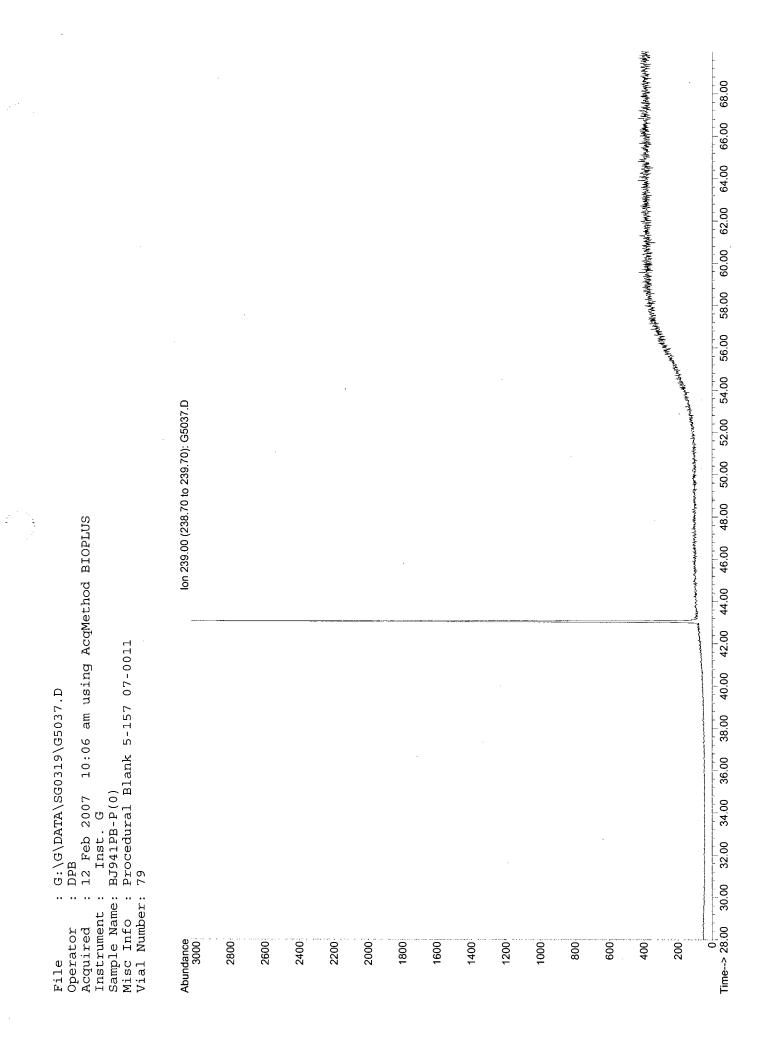


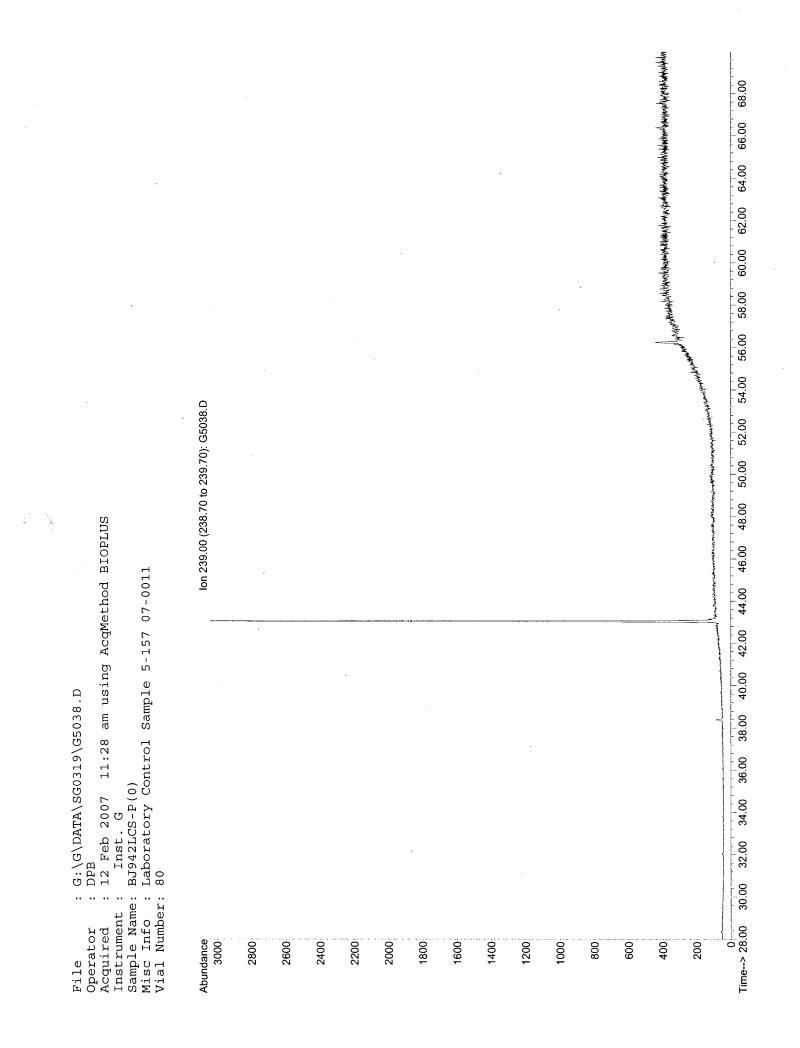


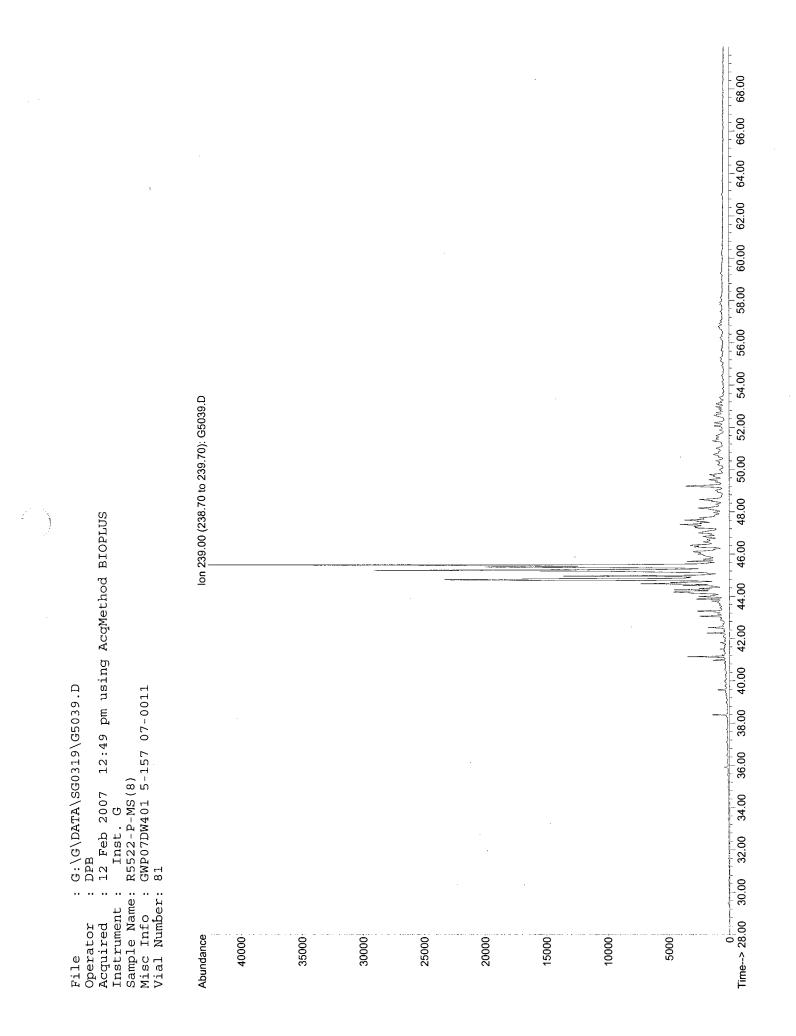


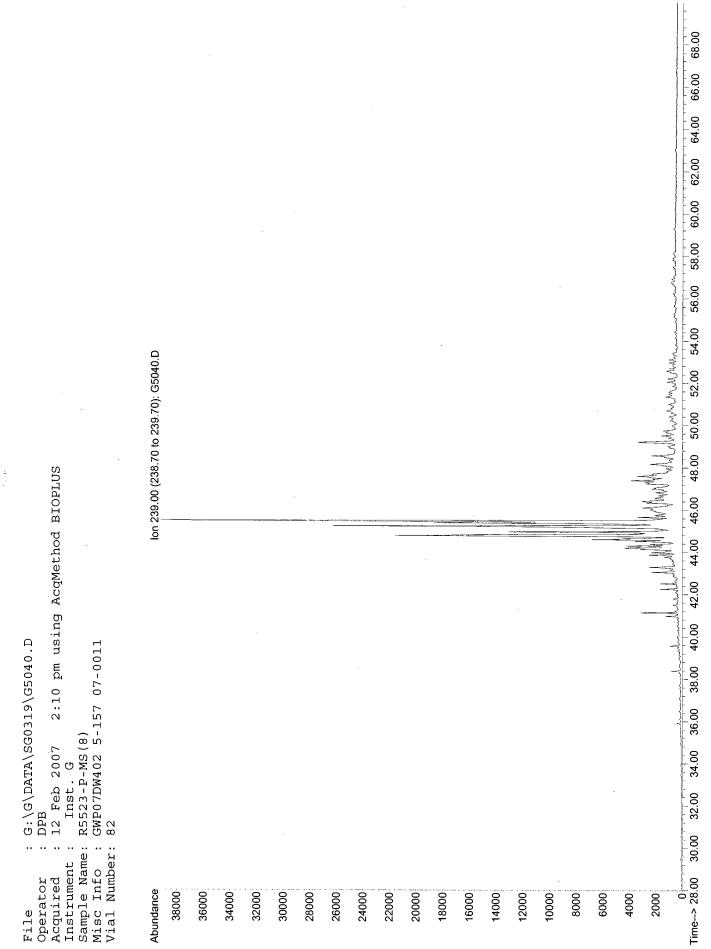
pm using AcqMethod BIOPLUS G:\G\DATA\SG0319\G5041.D DPB 12 Feb 2007 3:33 pm Inst. G R5524-P-MS(6) GWP07MW9 5-157 07-0011 83 8 Sample Name: Misc Info : Vial Number: ••• Instrument Operator Acquired File

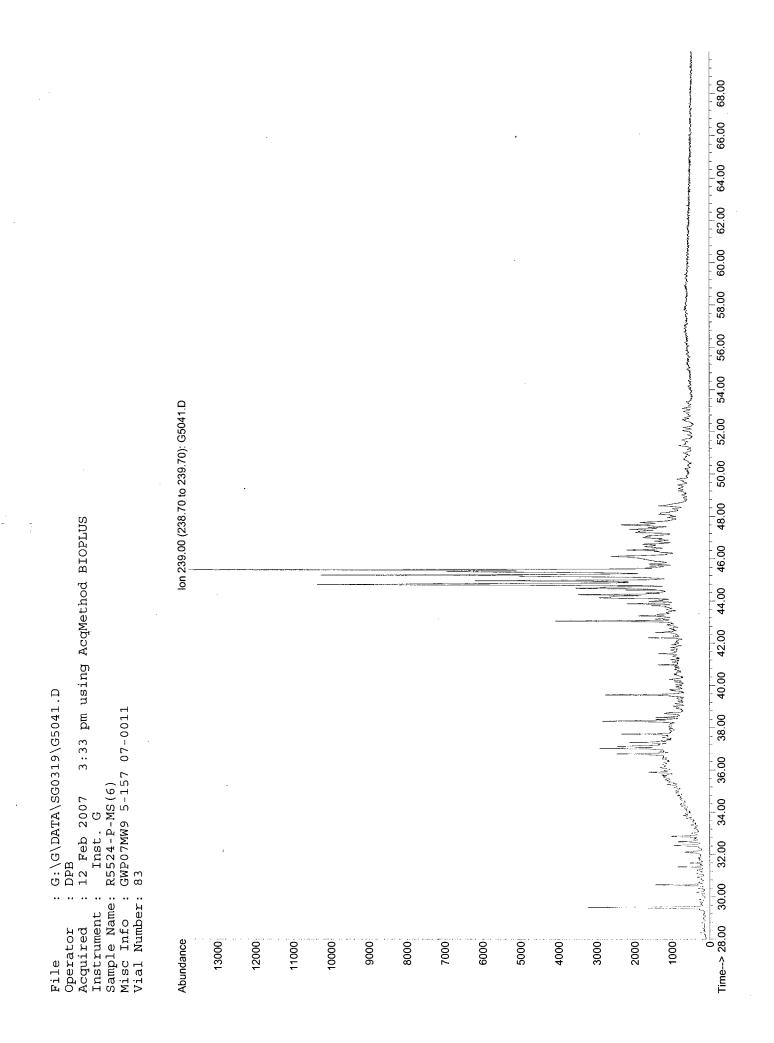












Data Qualifiers



Glossary of Data Qualifiers

Flag: Application:

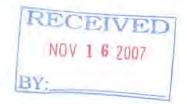
- B Analyte concentration found in the sample at a concentration <5x the level found in the procedural blank.
- D Dilution Run. Initial run outside linear range of instrument.
- E Estimate, result is greater than the highest concentration level in the calibration.
- H Surrogate diluted out. Used when surrogate recovery is affected by excessive dilution of the sample extract.
- J Analyte detected below the sample-specific Reporting Limit (RL).
- ME Significant Matrix Interference Estimated value.
- MI Significant Matrix Interference value could not be determined or estimated.
- n Quality Control (QC) value is outside the accuracy or precision Data Quality Objective (DQO), but meets the contingency criteria.
- N Quality Control (QC) value is outside the accuracy or precision Data Quality Objective (DQO)
- NA Not applicable
- T Holding Time (HT) exceeded.
- U Analyte not detected at 3:1 signal:noise ratio. The sample-specific method detection limit (MDL) reported.

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Tar Removal and Source Sampling Description Table

Sampling	Sample ID	Sample Type	Comments
Location	-		
DW-04	GWP07-DW4	Physical	Well located in Harbor Patrol parking lot
	GWP07-DW401	Forensics	
	GWP07-DW402	Forensics-duplicate	7
Station 1	GWP07-T01	Physical	Station east of play barn near blackberry
	GWP07-T01	Forensics	bushes, location with strong naphthalene odor
	GWP07-T02	Forensics	noticed during warmer temperatures. During
	GWP07-T03	Forensics-duplicate of T02	sampling PID reading of 541 units. Solidified tar
	GWP07-T04	Forensics	removed by Parks.
Station 2	GWP07-T05	Physical	Solidified tar seep previously covered with
	GWP07-T05	Forensics	gravel by Parks. Approx. 3' by 3', PID reading
	GWP07-T06	Forensics	around 7 units. Solidified tar removed by Parks.
	GWP07-T07	Forensics	7
	GWP07-T10	Physical	Soft vein of tar identified by Jodi and Parks
	GWP07-T10	Forensics	
Station 3	GWP07-T08	Physical	Station NW corner, roots of birch tree on
	GWP07-T08	Forensics	shoreline
	GWP07-T09	Forensics	7
Station 4	GWP07-T11	Physical	Aged solidified tar sample east of southerly play barn building. Tar between large pieces of
	GWP07-T11	Forensics	clinker.
Station 5	GWP07-T12	Physical	Aged solidified tar sample embedded with
	GWP07-T12	Forensics	gravel and some clinker. Slightly pliable, in
	GWP07-T13	Forensics	August/September during site visit with Marya
Station 6	GWP07-T14	Forensics	Possible solidified tar intermixed with dirt and gravel, faint naphthalene odor, small quantity collected (~1 oz).
Structure 1	GWP07-S01	Physical	Structure is a historical scrubber located in the
	GWP07-S01	Forensics	secured/fenced "Cracking Towers" area.
	GWP07-S02	Forensics	
Structure 2	GWP07-S03	Forensics	Structure is a historical scrubber located west of
			the play barn in fenced area, not locked. Only
			small quantity of tar available.
Structure 3	GWP07-S04	Physical	Structure is a historical scrubber located west of
	GWP07-S04	Forensics	structure 2 and the play barn in fenced area, not
MW-9	GWP07-MW9	Physical	Well located SW of play barn and SE of
	GWP07-MW9	Forensics	structures 2 and 3

ATTACHMENT 2D-6 Battelle November 2007 Data Report



Duxbury Operations 397 Washington Street Duxbury, Massachusetts 02332 Telephone 781-934-0571 Fax: 781-934-2124

Battelle

The Business of Innovation

November 15, 2007

Ms. Lisa Meoli Floyd/Snyder Two Union Square 601 Union Street, Suite 600 Seattle, WA 98101

Subject: Data Delivery - Gas Works Park

Dear Ms. Meoli:

Enclosed please find analytical data associated with the Gas Works Park project. The deliverable includes sample custody records, all analytical data tables (including QC data), GC chromatograms, PAH histograms, biomarker EICPs, and a data quality narratives associated with the data set. The narrative includes custody information, a summary of the processing and analysis methods, holding time information, and a discussion of issues related to quality control samples analyzed with the sample batch.

Please call me at (781) 952-5235 if you have any questions or you need additional information.

Sincerely,

Robert Lizotte,

Research Scientist

CS43	Normal	al TAT		- aga	10			Analytical Resources, Incorporated
ANI LIPHOUP I Swider	ev P	OC. COC. DOC	Stor	Date: 10]3	107	Present? V		4611 South 134th Place, Suite 100
2	Singal	2		No. of Coolers:	Temps:	er Aub)	206-695-6200 206-695-6201 (fax)
Ulent Froject Name:	5					Analysis Requested		Notes/Comments
cliphe Brolect #:	Samplers:	11 M. Kin		00	T	DILYZ		
Sample ID	Date	/ Time Matrix	No. Containers	Hes .	Thid	101		
p-WM	1 70 200	14:00 product	6.5)	7			LNAPI
		_						
					-			
Comments/Snerial Instructions	Balifordianad							
	HORC	e heude	(Senature) -	0	0	Relinquished by: (Signature)		Received by:
	Printed Name: N	deoli	Fringd Name:	1150.0		Printed Namo:		(organization) Printed Name:
	Floud	Swider	Company:			Company:		Company:
	Uate & Time:	taleal	Date & Time:		10/3/2	Date & Time:		Date & Time:

said services. The acceptance by the client of a proposal for services by ARI release ARI from any liability in excess thereof, not withstanding any provision to the contrary in any contract, purchase order or co-signed agreement between ARI and the Client.

Sample Retention Policy: All samples submitted to ARI will be appropriately discarded no sooner than 90 days after receipt or 60 days after submission of hardcopy data, whichever is longer, unless alternate retention schedules have been established by work-order or contract.

ARI Assigned Number: ARI Client Company	Turn-around Requested:	tequested:	Standard	rd	Date:	September 28, 2007	ber 28,	2007		Analytical Resources, Incorporated Analytical Chamiets and Consultation
Floyd/Snider		Phone: (2	Phone: (206) 292-2078	2-2078	Page		of			4611 South 134th Place, Suite 100
Jes					No. of Coolers:	0 P	Cooler Temps:	2.6)	1 ukwila, WA 98168 206-695-6200 206-695-6201 (fax)
COS-LCES	S							Analysis Requested		Notes/Comments
Client Project #: COS-LCES	Samplers:					-				
Sample ID	Date	Time	Matrix	No. Containers	Metho 8270	8270 by Extraction SPLP	8560	D× NMTP	-	
SB-2 S5 8-9.5	09/28/07		Soil				L			Please only use 2.5 oz and archive
SB-8 S5 9-10.5	09/28/07		Soil							the remainder for forensic analysis. Please only use 2.5 oz and archive
SB-13 25-4.0	09/28/07		Soil		>	>				the remainder for forsenic analysis.
						-	1	-		
SB-12A S3 5-6.5	09/28/07		Soil			+	+	-		Please archive total sample
							+			volume for forensic analysis.
						+	-			
							+			
							-			
Comments/Special Instructions	Pallocurried but									
Please see comments	(Signatore)	- lite	Hi.	Received by: (Signature)			Relir (Slor	Relinquished by: (Slaneture)		Received by
regarding sample volume. Hold partial volume for SB-2 and	Printed Name:	Meel		Printed Name:	ŀ		avint	Printed Name:		(signalue) Printed Name:
lction. B-12A	Company.	1 Sur	cler	Company.			E S	Company:		Company;
until turtner instruction.	Dates a time:	111	20	Date & Time:			Date	Date & Time:		Date & Taxe

c × Chain of Custody Record & Laborato

Limits of Liability: ARI will perform all requested services in accordance with appropriate methodology following ARI Standard Operating Procedures and the ARI Quality Assurance Program. This program meets standards for the industry. The total liability of ARI, its officers, agents, employees, or successors, arising out of or in connection with the requested services, shall not exceed the Invoiced amount for said services. The acceptance by the client of a proposal for services by ARI release ARI from any liability in excess thereof, not withstanding any provision to the contrary in any contract, purchase order or co-signed agreement between ARI and the Client.

Sample Retention Policy: Unless specified by workorder or contract, all water/soil samples submitted to ARI will be discarded or returned, no sooner than 90 days after receipt or 60 days after submission of hardcopy data, whichever is longer. Sediment samples submitted under PSDDA/PSEP/SMS protocol will be stored frozen for up to one year and then discarded.

ARI Assigned Number: ARI Client Company	Turn-around	Turn-around Requested:	Standard	ard	Date:	Septe	September 28, 2007	8, 2007	6	Analytical Resources, Incorporated
Floyd/Snider		Phone: (2	06) 29	Phone: (206) 292-2078	Page:	2	of		5	Analytical Chemists and Consultants 4611 South 134th Place, Suite 100
Jessi Massingale	singale				No. of Coolers:		Cooler Temps:	26)	Tukwila, WA 98168 206-695-6200 206-695-6201 (fax)
Gas Wol	Gas Works Park - COS (N	COS (N	E Corner)	1				Analysis Requested		Notas/Commante
Client Project #: Gas Works Park - COS	Samplers:				р			-+		S110111100-00000
Sample ID	Date	Time	Matrix	No. Containers	Metho 0728	Methoo 8260	SPLP Extraction (8270 by	D× NWTPł		
SB-3 S5 10-11.5	09/28/07		Soil		>	1				
SB-6 S2 1-2.0	09/28/07		Soil		>		5		1	
SB-10 S7 15-16.5	09/28/07		Soil		>		1			
GP-12 S2 23-24	09/28/07		Soil		>	>	T		+	
GP-1 12.5-13	09/28/07		Soil		>		1		+	
GP-11 14-14.5	09/28/07		Soil		1	>	T		+	
GP-12 S1 8-12	09/28/07		Soil			1	T		-	Plasen and the second
GP-9 7-8	09/28/07		Soil				T	-		and archive remainder.
						-	T			and archive remainder
Comments/Special Instructions	Retinutshed hv	-			H					
Please see comments	(Signature)	C. ILLY	ti.	Received by: (Signature)			R. S.	Reinquished by (Signature)		Received by: (Storature)
regarding sample volume.	U Si	Liet h		Printed Name:			ă	Printed Name:		Printed Name:
	1212	Firuel Sou	1. 1 C 1 C 1	Compuny			3	Company.		Company:
	La C La	11.21		Data & Time.			ő	Date & Time:		1010

The solution of the industry. The total liability of ARI, its officience with appropriate methodology following ARI Standard Operating Procedures and the ARI Quality Assurance Program. This program meets standards for the industry. The total liability of ARI, its officiens, agents, employees, or successors, anising out of or in connection with the requested services. Shall not exceed the Invoiced amount for said services. The acceptance DRI is program said services. The acceptance DRI is program said services. The acceptance DRI is program said services and the ARI Quality Assurance Program. This program said services. The acceptance DRI is program said services and the client of a proposal for services by ARI release ARI from any liability in excess thereof, not withstanding any provision to the contrary in any contract, purchase order or co-signed agreement between ARI and the Client.

Sample Retention Policy: Unless specified by workorder or contract, all water/soil samples submitted to ARI will be discarded or returned, no sooner than 90 days after receipt or 60 days after submission of hardcopy data, whichever is longer. Sediment samples submitted under PSDDA/PSEP/SMS protocol will be stored frozen for up to one year and then discarded.

The Business of Innovation

Data Report

Gas Works Park November 2007

Prepared For:

Lisa Meoli Floyd/Snider Two Union Square 601 Union Street Seattle, WA 98101

Prepared By:

Battelle 397 Washington Street Duxbury, Massachusetts 02332 Chain of Custody

Battelle The Business of Innovation

ShpNo SHP-071009-02

Battelle Project No: OP47854

Commis	Description Discourses	
SETTION:	Receipt Form	
and the second second		

and the second	and the second s	and and and	Approved:	
Project Number:		Client: ARI / Floyd-Sni	der	
Received by:	Seyfert, Jeannine	Date/Time Received:	Tuesday, October 09, 2007	12:00 AM
No. of Shipping Con	tainers: 1			
SHIPMENT				
Method of Delivery:	Commercial Carrier	Tracking Number:	1Z-832-695-01-4511-8864	
COC Forms:	Shipped with samples	No Forms		
Cooler(s)/Box				
Cntr Type 1 of 1 Cardboard B	Tracking No.	Seal Seal Condition	and the second s	Temp C Smps
Samples	07 12-002-030-01-4011-0004	Tape Intact	Intact	0.6 7
Sample Labels:				
ample Ladels:		agree with COC forms		
	Discrepancies	(see Sample Custody Corre	ective Action Form)	
Container Seals:	🗌 Tape 🔲 Cu	istody Seals 🗌 Other Sea	le (See cample Loc)	
	Seals intact fo	r each shipping container	is (see sample Log)	
		See sample log for impacted	d samples)	
Condition of Samples:	Sample contai	ners intact		
	Sample contai	ners broken/leaking (See Cu	ustody Corrective Action Fo	rm)
emperature upon reco Note: If temperature up	eipt (°C): 0.6 Te oon receipt differs from required co	mperature Blank used	Yes 🔽 No	
	on receipt aggers from required co	mattions, see sample log co	omment field)	
amples Acidified:	🗌 Yes 🗌 No	Unknown		
nitial pH 5-9?:	🗌 Yes 🗌 No	NA NA		
no, individual sample	adjustments on the Auxiliary Samp	ole Receipt Form		
otal Residual Chlorine		✓ NA		
yes, individual sample	adjustments on the Auxiliary Sam	ple Receipt Form		
lead Snace <1% in sam	nples for water VOC analysis:			
dividual sample deviat	ions noted on sample log	Yes No V NA	A.	
	ions noted on sumple log			
amples Containers:				
amples returned in PC-g	grade jars: 🖌 Yes 🗌 No	Unknown /Lot No.: U	InKnown	
orage Location:	Chem North: Freezer - F0002			
amples logged in by:	Seyfert, Jeannine	(maiking BDO ID	os Assigned: Q0540 - Q	
pproved By:	Brackett, Roxanne			9/2007 12:00 AM
	Brackett, Koxanne		Approved On: 11/12	2/2007 12:
uthorized By:			Authorized On:	

Battelle The Business of Innovation

ShpNo: SHP-071009-02

Battelle Project No: OP47854

COC Client:	ARI / Floyd-S	Snider		
COC Project:	Gasworks Pa	rk		
COC Date:	10/9/2007 1:	42:00 PM		
		ion of Problem:	Expla	nation:
Sample Container Integrity	Sample leakir	IG	residue well as it was on the	e Q0540 (MW-9) had some oil e on the outside of the container as s on the inner side of the bubble wrap wrapped in. The oil residue was only inside of the bubble wrap and did uch any of the other samples.
Temperature and Preservation	Receipt tempe	erature outside of acceptability	The co	oler temperature upon receipt was of the 4+/- 2 degree range of
Documentation	of project mana	ger notification		
Sample Cu	stodian	Seyfert, Jeannine	Date:	10/9/2007 1:58:00 PM
Laboratory	Manager:	Thorn, Jonathan	Date:	11/6/2007 7:33:00 AM
Project Ma		Krahforst, Kerylynn	Date:	11/12/2007 12:00:00 P
Documentation of	of client notifica	tion (should be completed by project	manager wit	thin 24 hrs):
On 16-0	ct-07 I cor	ntacted Meoli, Lisa	at Flo	oyde Snider

Date this form was received back to the custodian:

The Business of Innovation Battelle

ShpNo SHP-071009-02

OP47854 Battelle Project No:

Sample Receipt Form Details

Approved: 🗸

Client: ARI / Floyd-Snider Seyfert, Jeannine Project Number: Received by:

Date/Time Received: Tuesday, October 09, 2007 12:00 AM

No. of Shipping Containers: 1

- and the second	Collection Date:	Login Date:	Ctre. Matriv.	Towner						
20540 MW-9			CUS. MGUIA.	IEMP: PH: IKC: VOC:	11	C: NO	:: Stored In:	Loc:	No: Comments:	Iments:
SP 7 SE 8 DE	10/03/07 0:00	10/09/07 13:47	1 NAPL	0.6 N	AN	A NA	F0002 (Walk-in)	BIN	87 07-2	7-21114-1 SA3A
SR-8 S5 0-10 5	00:0 /0// 1/60	10/09/07 13:47	1 SOIL	0.6 N	AN	A NA	F0002 (Walk-in)	BIN	87 07-2	07-20838-I R81A
SB-13 2 5 4 0	00:0 /0/01/60	10/09/07 13:48	1 SOIL	0.6 N	AN	A NA	F0002 (Walk-in)	BIN	87 07-2	07-20839-I R81B
SB-124 S3 6 6 6	00:0 /0/07/60	10/09/07 13:49	1 SOIL	0.6 N	AN	AN NA	F0002 (Walk-in)	BIN	87 07-2	07-20840-I R81C
GP-12 S2 23-24	00:0 /0/07/60	10/09/07 13:50	1 SOIL	0.6 N	AN	A NA	F0002 (Walk-in)	BIN	87 07-2	07-20842-LR81F
00546 GP-0 7.8	00:0 /0/07/60	10/09/07 13:52	1 SOIL	0.6 N	AN	A NA	F0002 (Walk-in)	BIN	87 07-2	07-20828-I R80D
2	03/18/07 0:00	10/09/07 13:53	1 SOIL	0.6 N	NA N	NA NA	F0002 (Walk-in)	BIN	0 70 79	07 20022 L D0011

SUBCONTRACTOR ANALYSIS REQUEST CUSTODY TRANSFER 10/04/07



Laboratory: Battelle Laboratories Duxbury ARI Client: Floyd-Snider Lab Contact: Kerylynn Krahforst Lab Address: 397 Washington Street Duxbury, MA 02332 Phone: Fax:

Project ID: Gasworks ARI PM: Sue Dunnihoo Phone: 206-695-6207 Fax: 206-695-6201

Analytical Protocol: In-house Special Instructions:

Requested Turn Around: 10/17/07 Fax Results (Y/N): Y

Limits of Liability. Subcontractor is expected to perform all requested services in accordance with appropriate methodology following Standard Operating Procedures that meet standards for the industry. The total liability of ARI, its officers, agents, employees, or sucessors, arising out of or in connection with the requested services, shall not exceed the negotiated amount for said services. The agreement by the Subcontractor to perform services requested by ARI releases ARI from any liability in excess thereof, not withstanding any provision to the contrary in any contract, purchase order or co-signed agreement between ARI and the Subcontractor.

ARI ID	Client ID/ Add'l ID	141	Sampled	Matrix	Bottles	Analyses	Battelle ID
07-21114-LS43A	MW-9		10/03/07	Product	1	WILDCARD	QC540
Special Instruc	tions: None						01-01-0

Carrier UPS Religiuished by	Airbill 12832695	01 4511 8464	Date 10/8/7	1
D'ZIR.	Company Refelle Dichard	Date 10/8/12 Date	Time 1600	111
J JI	Subcontractor Custody P Page 1 of	Form - LS43	13.00	

SUBCONTRACTOR ANALYSIS REQUEST CUSTODY TRANSFER 10/04/07



Laboratory: Battelle Laboratories Duxbury ARI Client: Floyd-Snider Lab Contact: Kerylynn Krahforst Lab Address: 397 Washington Street Duxbury, MA 02332 Phone: Fax:

Project ID: COS-LCES ARI PM: Sue Dunnihoo Phone: 206-695-6207 Fax: 206-695-6201

Analytical Protocol: In-house Special Instructions:

Requested Turn Around: 10/17/07 Fax Results (Y/N): Yes

Limits of Liability. Subcontractor is expected to perform all requested services in accordance with appropriate methodology following Standard Operating Procedures that meet standards for the industry. The total liability of ARI, its officers, agents, employees, or sucessors, arising out of or in connection with the requested services, shall not exceed the negotiated amount for said services. The agreement by the Subcontractor to perform services requested by ARI releases ARI from any liability in excess thereof, not withstanding any provision to the contrary in any contract, purchase order or co-signed agreement between ARI and the Subcontractor.

Client ID/ Add'l ID	Sampled	Matrix	Bottles	Analyses	Enthelle ID
SB-2 S5 8-9.5	09/17/07	Soil	1	WILDCARD	Q0541
tions: None					
SB-8 S5 9-10.5	09/18/07	Soil	1	WILDCARD	R0592
tions: None					
SB-13 2.5-4.0	09/20/07	Soil	1	WILDCARD	Q0543
tions: None					40570.
SB-12A S3 5-6.5	09/20/07	Soil		WILDCARD	20544
ions: None					
	Add'1 ID SB-2 S5 8-9.5 tions: None SB-8 S5 9-10.5 tions: None SB-13 2.5-4.0 tions: None SB-12A S3 5-6.5	Add'1 ID Sampled SB-2 S5 8-9.5 09/17/07 tions: None 09/18/07 SB-8 S5 9-10.5 09/18/07 tions: None 09/20/07 SB-13 2.5-4.0 09/20/07 tions: None SB-12A S3 5-6.5 SB-12A S3 5-6.5 09/20/07	Add'1 ID Sampled Matrix SB-2 S5 8-9.5 09/17/07 Soil tions: None 09/18/07 Soil tions: None 09/20/07 Soil	Add'1 ID Sampled Matrix Bottles SB-2 S5 8-9.5 09/17/07 Soil 1 tions: None 09/18/07 Soil 1 sB-8 S5 9-10.5 09/18/07 Soil 1 tions: None 09/20/07 Soil 1 tions: None 09/20/07 Soil 1 sB-13 2.5-4.0 09/20/07 Soil 1 tions: None 09/20/07 Soil 1 sB-12A S3 5-6.5 09/20/07 Soil 1 tions: None 1 1 1	Add'1 ID Sampled Matrix Bottles Analyses SB-2 S5 8-9.5 09/17/07 Soil 1 WILDCARD tions: None 09/18/07 Soil 1 WILDCARD tions: None 09/20/07 Soil 1 WILDCARD sB-12A S3 5-6.5 09/20/07 Soil 1 WILDCARD tions: None 1 WILDCARD 1

Carrier UPS	Airbill		Date
Received by flanture fight	Company Company Battelle Duxbery	Date 10/8/7 Date 0/9/7	Time 13:00
U	Subcontractor Custody F Page 1 of	Sorm - LR81 1	10 m 11 m 16 m

SUBCONTRACTOR ANALYSIS REQUEST CUSTODY TRANSFER 10/04/07

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Laboratory: Battelle Laboratories Duxbury ARI Client: Floyd-Snider Lab Contact: Kerylynn Krahforst Lab Address: 397 Washington Street Duxbury, MA 02332 Phone: Fax:

Project ID: Gas Works Park-COS (NE Corner) ARI PM: Sue Dunnihoo Phone: 206-695-6207 Fax: 206-695-6201

Analytical Protocol: In-house Special Instructions:

Requested Turn Around: 10/17/07 Fax Results (Y/N): Yes

Limits of Liability. Subcontractor is expected to perform all requested services in accordance with appropriate methodology following Standard Operating Procedures that meet standards for the industry. The total liability of ARI, its officers, agents, employees, or sucessors, arising out of or in connection with the requested services, shall not exceed the negotiated amount for said services. The agreement by the Subcontractor to perform services requested by ARI releases ARI from any liability in excess thereof, not withstanding any provision to the contrary in any contract, purchase order or co-signed agreement between ARI and the Subcontractor.

ARI ID	Client ID/ Add'l ID	Sampled	Matrix	Bottles	Analyses	
07-20828-LR80	D GP-12 S2 23-24	09/20/07	Soil	1	WILDCARD	Q0545
Special Instr	uctions: None					
07-20832-LR80	H GP-9 7-8	09/18/07	Soil	1	WILDCARD	Q0.546
Special Instr	uctions: None					4-0-10

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SHC Data and Chromatograms

SHC and TPH – SEDIMENT QA/QC SUMMARY Batch 07-0259

PROJECT:	Floyd/Snyder – Gas Works Park
PARAMETER:	Saturated Hydrocarbons (SHC) and Total Petroleum Hydrocarbons (TPH)
LABORATORY:	Battelle, Duxbury, MA
MATRIX:	Sediment
SAMPLE CUSTODY:	Six soil samples and 1 NAPL were received at Battelle Duxbury Operations (BDO) Laboratory on 10/9/07. Upon Receipt of the samples, the temperature of the cooler was taken and the samples were logged into the laboratory and given unique IDs. The temperature of the cooler upon receipt was slightly below the acceptable range ($4^{\circ}C \pm 2^{\circ}$) at 0.6°C. Also, it was noted that sample Q0540 (MW-9) had some oil residue on the outside of the container as well as the inner side of the bubble wrap. The oil residue was only on the inside of the bubble wrap and did not touch any of the other samples. The client was notified on 10/16/07. The laboratory was instructed to proceed with the analysis. Samples were stored in an access-limited walk-in refrigerator at 4°C until sample preparation could begin.

	Reference Method	Method Blank	Surrogate Recovery	LCS Recovery	MS Recovery	MS/MSD Precision	Control Oil % Diff
SHC and TPH	General NS&T	<5xMDL	40-120% Recovery	40-120% Recovery	40-120% Recovery	≤30% RPD	PD,30% for 90% of the analytes
					MS target spike must be >5x background		

METHOD:

Soil samples were extracted following general NS&T methods. Approximately 5-8 g of sample was spiked with SHC and PAH surrogates and serial extracted three times with dichloromethane using orbital shaker table techniques. The combined extracts were dried over anhydrous sodium sulfate and concentrated by Kuderna-Danish and nitrogen evaporation techniques. The sample extracts were split in half: one-half of the extract was removed for archiving; the other half was processed through an alumina gravity column to isolate the hydrocarbon fractions of interest. The weights of the resulting extracts were determined gravimetrically. The extracts were concentrated to 1 mL, split, and spiked with internal standard (IS). The pre-injection volume and/or extract split were adjusted to 5mg/mL. One extract was submitted for PAH and the second extract was submitted for SHC and TPH analysis.

SHC and TPH were measured by gas chromatography with flame ionization detection (GC/FID). An initial calibration consisting of target analytes was completed prior to analysis to demonstrate the linear range of the analysis. Calibration verification was performed at the beginning and end of each 12 hour period in which samples were analyzed. Concentrations of SHC and TPH were calculated by the internal standard method. Normal alkanes were quantified using the average RF generated from the initial calibration. TPH concentrations were quantified suing the average RF of nC8 through nC40. Isoprenoid hydrocarbon concentrations were quantified using the average RF of nc8 through nC40. Isoprenoid hydrocarbon concentrations were quantified using the average RF of nc8 through nC40.

HOLDING TIMES: Samples were prepared for analysis in three analytical batches and were extracted within 30 days of sample collection analyzed within 40 days of extraction.

Batch	Extraction Date	Analysis Date
07-0259	10/15/2007	10/18/2007 - 10/19/2007

SHC and TPH – SEDIMENT QA/QC SUMMARY Batch 07-0259

PROCEDURAL BLANK (PB):	A procedural blank (PB) was prepared with each analytical batch. Blanks were analyzed to ensure the sample extraction and analysis methods were free of contamination.
	07-0259 – No exceedences noted.
	Comments – None.
LABORATORY CONTROL SAMPLE (LCS):	A laboratory control sample (LCS) was prepared with each analytical batch. The percent recoveries of target analytes were calculated to measure data quality in terms of accuracy. 07-0259 – No exceedences noted.
	Comments – None.
SURROGATE RECOVERY:	Two surrogate compounds were added prior to extraction, including o-terphenyl and 5a- androstane. The recovery of the surrogate compound was calculated to measure data quality
	in terms of accuracy (extraction efficiency).
	07-0259 – No exceedences noted.
	Comments – None.
CONTROL OIL:	A control oil (North Slope Crude) was prepared with the analytical batch. The percent difference (PD) between the measured value and the target value was calculated to measure data quality in terms of accuracy.
	07-0259 – No exceedences noted.
	Comments – None.
CALIBRATIONS:	The GC/FID is calibrated with a minimum 6 local sector is a local sector.
	The GC/FID is calibrated with a minimum 5 level curve for all compounds. The percent relative standard deviation (% RSD) between RF for the individual target analytes must be $\leq 25\%$, and the mean RSD of all target analytes must be $\leq 20\%$. Each batch of samples analyzed is bracketed by a continuing calibration verification (CCV) sample, run at a
	frequency of minimally every 12 hours. The PD between the true value and the CCV should be $\leq 25\%$ for individual analytes. Additionally an initial calibration check (ICC) sample is run immediately after each initial calibration. The PD between the ICC and the initial calibration should be $\leq 25\%$.
	07-0259 – No exceedences noted.
	Comments - None

The Business of Innovation

Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	SB-2 S5 8-9.5	GP-12 S2 23-24	
Battelle ID	Q0541-P	20515 2	
Sample Type		Q0545-P	
Collection Date	SA	SA	
Extraction Date	09/17/07	09/20/07	
Analysis Date	10/15/07	10/15/07	
Analysis Date	10/18/07	10/19/07	
Analytical Instrument	FID	FID	
% Moisture	18.05	26.35	
% Lipid	NA	NA	
Matrix	SOIL	SOIL	
Sample Size	4.59	3.93	
Size Unit-Basis	G_DRY	G DRY	
Units	UG/G_DRY	UG/G_DRY	
n-Nonane	U		
n-Decane	Ű	ů.	
n-Undecane		U	
n-Dodecane	U	U	
n-Tridecane	U	U	
	U	U	
Isoprenoid RRT 1380	U	U	
n-Tetradecane	U	Ū	
Isoprenoid RRT 1470	U	U	
n-Pentadecane	165.07	9.04	
n-Hexadecane	86.55	28.62	
Norpristane (1650)	U	U	
n-Heptadecane	Ŭ	Ŭ	
Pristane	213.07	ŭ	
n-Octadecane	228.64		
Phytane	180.99	82.21	
n-Nonadecane		22.17	
n-Eicosane	4.97	1.05 J	
n-Heneicosane	U	U	
	U	9.78	
n-Docosane	U	U	
n-Tricosane	U	U	
n-Tetracosane	U	U	
-Pentacosane	U	U	
n-Hexacosane	U	U	
n-Heptacosane	U	Ŭ	
-Octacosane	U	- ŭ	
n-Nonacosane	Ũ	11.65	
-Triacontane	ŭ		
-Hentriacontane	Ŭ	U	
-Dotriacontane	U	0 U	
-Tritriacontane	Ŭ	2.18	
-Tetratriacontane		6.11	
-Pentatriacontane	U	3.08	
-Hexatriacontane	U	U	
	U	U	
-Heptatriacontane	U	U	
-Octatriacontane	U	U	
-Nonatriacontane	U	U	
-Tetracontane	Ŭ	Ű	
PH(total)	19288.58	2228.21	
urrogate Recoveries (%)			
Terphenyl	440		
a-androstane	113	96	

The Business of Innovation

Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	Procedural Blank	
Battelle ID	BL033PB-P	
Sample Type	PB	
Collection Date		
Extraction Date	10/15/07	
Analysis Date	10/15/07	
Analytical Instrument	10/18/07	
	FID	
% Moisture	22.07	
% Lipid	NA	
Matrix	SEDIMENT	
Sample Size	11.03	
Size Unit-Basis	G_DRY	
Units	UG/G_DRY	
n-Nonane	U	
n-Decane	U	
n-Undecane	Ũ	
n-Dodecane	Ŭ	
n-Tridecane	ŭ	
Isoprenoid RRT 1380	Ü	
n-Tetradecane	U	
Isoprenoid RRT 1470	0	
n-Pentadecane	U U	
n-Hexadecane		
Norpristane (1650)	U.	
n-Heptadecane	U	
Pristane	U	
n-Octadecane	U	
Phytane	U	
n-Nonadecane	U	
n-Eicosane	U	
n-Heneicosane	υ	
	U	
n-Docosane	U	
n-Tricosane	U	
n-Tetracosane	Ú	
n-Pentacosane	U	
n-Hexacosane	U	
n-Heptacosane	U	
n-Octacosane	U	
n-Nonacosane	U	
n-Triacontane	U	
n-Hentriacontane	U	
n-Dotriacontane	U	
n-Tritriacontane	U	
n-Tetratriacontane	U	
n-Pentatriacontane	Ŭ	
n-Hexatriacontane	Ŭ	
n-Heptatriacontane	Ŭ	
n-Octatriacontane	Ŭ	
n-Nonatriacontane		
n-Tetracontane	ũ	
TPH(total)		
Surrogate Recoveries (%)		
O-Terphenyl	91	
5a-androstane	91	

O-Terphenyl 5a-androstane

94

The Business of Innovation

Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	060208-03: Sand, White Quartz, -50+70					
Battelle ID	BL034LCS-P					
Sample Type	BL034LCS-P LCS					
Collection Date	10/15/07					
Extraction Date	10/15/07					
Analysis Date	10/18/07					
Analytical Instrument	FID					
% Moisture	NA					
% Lipid	NA					
Matrix	SEDIMENT					
Sample Size	NA					
Size Unit-Basis	NA					
Units	UG		Target 9	& Recovery	Qualifier	
n-Nonane	70.5		100.00	44		
n-Decane	79.72		100.00	71 80		
n-Undecane	74.9		100.00	80 75		
n-Dodecane	79		100.00	75 79		
n-Tridecane	78.16		100.00	79		
soprenoid RRT 1380	75:16	U	100.00	10		
n-Tetradecane	79.07	-	100.00	79		
soprenoid RRT 1470	10.01	U	100.00	19		
-Pentadecane	81.89		100.00	82		
-Hexadecane	82.76		100.00	83		
lorpristane (1650)		U		00		
-Heptadecane	81.82	2	100.00	82		
Pristane	83.15		100.00	83		
-Octadecane	83.37		100.00	83		
hytane	82.3		100.00	82		
-Nonadecane	83.61		100.00	84		
-Eicosane	84.32		100.00	84		
-Heneicosane	83.54		100.00	84		
-Docosane	86.84		100.00	87		
-Tricosane -Tetracosane	84.72		100.00	85		
Pentacosane	85.21		100.00	85		
Hexacosane	86.59		100.00	87		
Heptacosane	84.71		100.00	85		
Octacosane	85.55		100.00	86		
Nonacosane	82.19		100.00	82		
Triacontane	87.14 82.99		100.00	87		
Hentriacontane	82.99		100.00	83		
Dotriacontane	81.33		100.00	81		
Tritriacontane	81.16		100.00	83		
Tetratriacontane	78.57		100.00	81 79		
Pentatriacontane	79.91		100.00	80		
Hexatriacontane	71.72		100.00	72		
Heptatriacontane	67.74		100.00	68		
Octatriacontane	65.06		100.00	65		
Nonatriacontane	58.22		100.00	58		
Tetracontane	53.38		100.00	53		
PH(total)		J	199100			

Surrogate Recoveries (%)

O-Terphenyl	82
5a-androslane	86

Battelle The Business of Innovation

Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	GO98: North Slope Crude				
Battelle ID	BL041NSC-P				
Sample Type					
Collection Date	NSC				
Extraction Date	10/17/07				
Analysis Date	10/17/07				
Analytical Instrument	10/18/07				
% Moisture	FID				
% Lipid	NA.				
Matrix	NA				
Sample Size	OIL				
Size Unit-Basis	5.04				
Units	G_OIL	Sec. 1			
Units	UG/G_OIL	Target %	Difference	Qualifier	_
n-Nonane	5.03	4.67	7.7		
n-Decane	4.79	4.95	3.3		
n-Undecane	4.31	4.51	4.4		
n-Dodecane	4.17	4.58	8.9		
n-Tridecane	4.02	4.19	4.0		
Isoprenoid RRT 1380	0.9	0.96	6.4		
n-Tetradecane	3.85	3.92	1.8		
soprenoid RRT 1470	1.5	1.53	2.1		
n-Pentadecane	3.86	3.99	3.3		
n-Hexadecane	3.34	3.64	8.2		
Norpristane (1650)	1.16	1.14	1.6		
n-Heptadecane	3.11	3.08	1.0		
Pristane	2.23	2.28	2.2		
1-Octadecane	2.88	2.80	3.0		
Phytane	1.4	1.66	15.7		
n-Nonadecane	2.56	2.54	0.8		
I-Eicosane	2.53	2.50	1.1		
I-Heneicosane	2.24	2.42	7.4		
-Docosane	2.21	2.25	1.9		
-Tricosane	1.98	2.05	3.4		
-Tetracosane	1.89	1.95	3.0		
-Pentacosane	1.71	1.80	4.8		
-Hexacosane	1.58	1.64	3.6		
-Heptacosane	1.18	1.23	4.1		
-Octacosane	0.94	1.00	6.4		
-Nonacosane	0.8	0.87	8.3		
-Triacontane	0.67	0.67	0.1		
Hentriacontane	0.66	0.61	8.8		
Dotriacontane	0.42	0.47	9.9		
Tritriacontane	0.43	0.40	7.8		
Tetratriacontane	0.34	0.37	8.5		
Pentatriacontane	0.4	0.38	5.8		
Hexatriacontane	0.23 J	0.24	2.4		
Heptatriacontane	0.2 J	0.21	4.8		
Octatriacontane	0.21 J	0.21	2.1		
Nonatriacontane	0.16 J	0.15	4.0		
Tetracontane PH(total)	0.15 J	0.16	7.2		
			1.6		

Surrogate Recoveries (%)

O-Terphenyl	95
5a-androstane	100



: 10-18-2007 01:09:38 PM using AcqMethod TPH.M : Inst. N : BL044SOL-P(0) : F:\N\DATA\SN0256.SEC\N5789.D : MM File : Operator : Acquired : Instrument : Sample Name:



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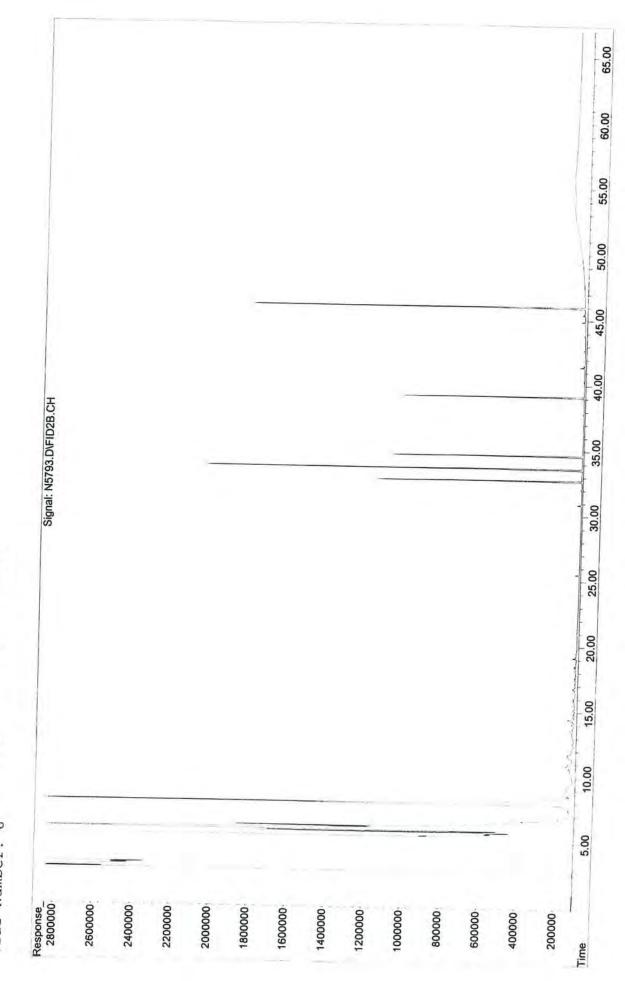
	TPH.M	
	AcqMethod TPH.M	
F:\N\DATA\SN0256.SEC\N5791.D MM	10-18-2007 02:29:27 PM using Inst. N	BL045SOL-P(0) Solvent blank. 5-202 07-0259 5
File : Operator :	Acquired : Instrument :	Sample Name: Misc Info : Vial Number:

Response	3400000	3200000	000000	2800000	2600000	2400000	2200000	2000000	1800000	1600000	1400000	1200000	100000	800000	600000	400000	200000
Signal: N																	
Signal: N5791.D/FID2B.CH																	
CH																	
										7							



	AcqMethod
193.D	using
SNI	Md
56.SEC\N5793	03:49:20
\SN0256	03:
F:\N\DATA\: MM	10-18-2007 Inst N

	TPH.M	
	AcqMethod	6
C\N5793.D	using	07-025
SE	:20 PM	5-202
SN0256	03:49	FID(4) Blank
F:\N\DATA\SN0256. MM	10-18-2007 Inst. N	
File : Operator :	cquired nstrumen	Sample Name: Misc Info : Vial Number:





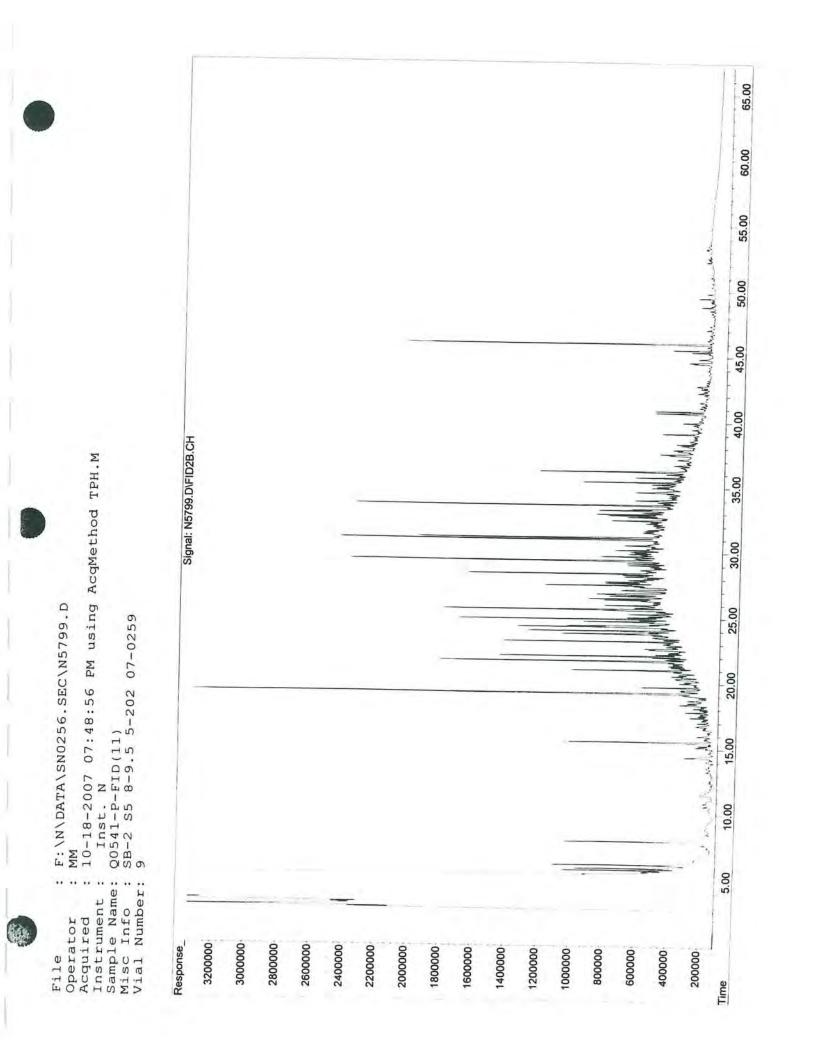
10-18-2007 05:09:34 PM using AcqMethod TPH.M Inst. N BL034LCS-P-FID(4) Laboratory Control Sample 5-202 07-0259 F:\N\DATA\SN0256.SEC\N5795.D MM 5 File : F Operator : M Acquired : 1 Instrument : Sample Name: E Misc Info : 1 Vial Number: 7

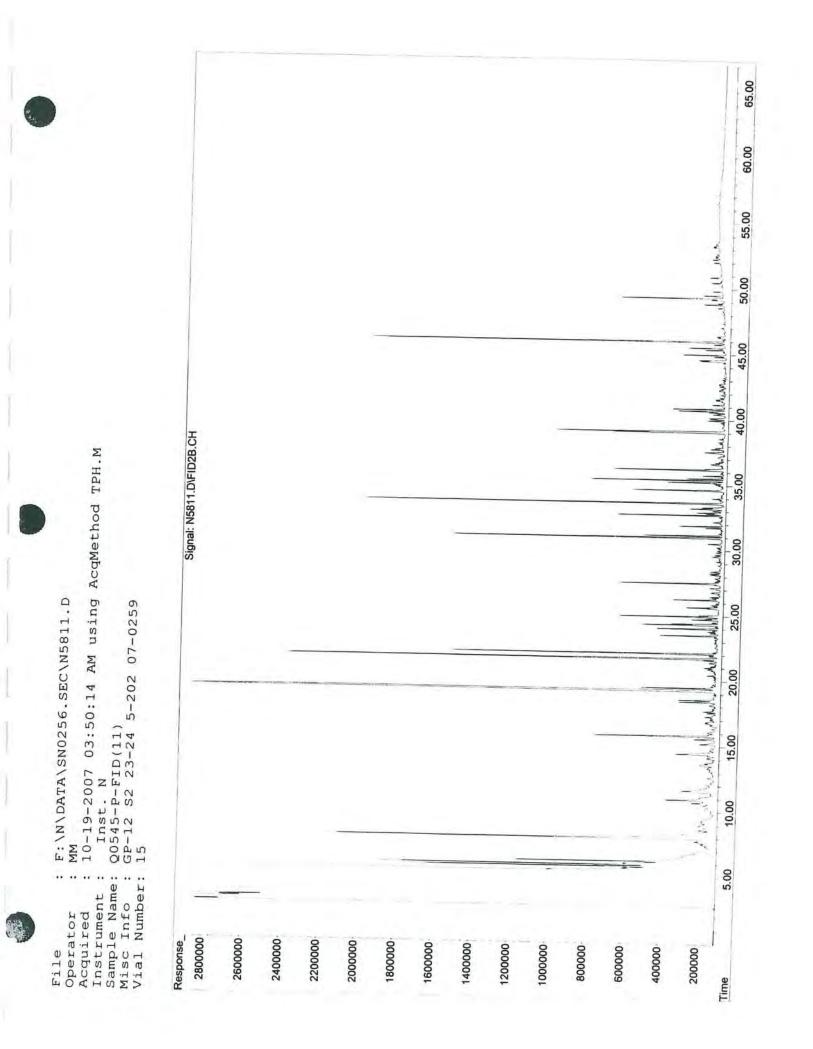
0.00 25.00	Signal: N5795.D/FID2B.CH		1111111					30.00 35.00 40.00
2				 				



10-18-2007 06:29:05 PM using AcqMethod TPH.M Inst. N BL041NSC-P(0) North Slope Crude 07-0259 8 F:\N\DATA\SN0256.SEC\N5797.D MM File : 1 Operator : 1 Acquired : 1 Instrument : Sample Name: F Misc Info : 1 Vial Number: 6

					 		and all the property of the function of the second s		
signal: N5/97.D/FID2B.CH							LUNK AN WAVAN MUNAMURINA ALLA	non-security and a start for a line of the second	





SHC and TPH - SEDIMENT QA/QC SUMMARY Batch 07-0264

PROJECT PARAME LABORAT MATRIX: SAMPLE	TER: FORY:	Saturated H Battelle, Du Non-aqueou Six soil sam Laboratory of taken and th temperature 2°) at 0.6°C. outside of th only on the i client was no analysis. Sat	er – Gas Works ydrocarbons (SF xbury, MA is phase liquid (N ples and 1 NAP on 10/9/07. Upo e samples were 1 of the cooler up Also, it was no e container as w nside of the bub otified on 10/16/ mples were store aration could beg	IC) and Total P NAPL) L were received on Receipt of the logged into the on receipt was s ted that sample ell as the inner ble wrap and di 07. The labora ed in an access-	at Battelle Du e samples, the t laboratory and slightly below t Q0540 (MW-9 side of the bubl d not touch any atory was instru	xbury Operatio emperature of t given unique II he acceptable r had some oil ble wrap. The of the other sa	ns (BDO) the cooler was Ds. The ange $(4^{\circ}C \pm$ residue on the oil residue was mples. The with the
	Reference Method	Method Blank	Surrogate Recovery	LCS Recovery	MS Recovery	MS/MSD Precision	Control Oil % Diff
SHC and	General	<5xMDL	40-120%	40-120%	40-120%	≤30% RPD	PD,30% for

Recovery

MS	target spike
mus	t be >5x
back	ground

Recovery

90% of the analytes

METHOD:

TPH

NS&T

The NAPL sample was extracted following general NS&T methods. Approximately 50 mg of oil was weighed and diluted with 10mL of hexane. A portion of the extract was removed and spiked with SIS and IS. One extract was submitted for PAH and the second extract was submitted for SHC and TPH analysis. NAPL sample data is reported on an oil weight basis.

Recovery

SHC and TPH were measured by gas chromatography with flame ionization detection (GC/FID). An initial calibration consisting of target analytes was completed prior to analysis to demonstrate the linear range of the analysis. Calibration verification was performed at the beginning and end of each 12 hour period in which samples were analyzed. Concentrations of SHC and TPH were calculated by the internal standard method. Normal alkanes were quantified using the average RF generated from the initial calibration. TPH concentrations were quantified suing the average RF of nC8 through nC40. Isoprenoid hydrocarbon concentrations were quantified using the average RF of the n-alkanes immediately preceding and immediately following each target isoprenoid hydrocarbon.

HOLDING TIMES:

Samples were prepared for analysis in three analytical batches and were extracted within 15 days of sample collection analyzed within 40 days of extraction.

Batch	Extraction Date	Analysis Date
07-0264	10/18/2007	10/19/2007

SHC and TPH – SEDIMENT QA/QC SUMMARY Batch 07-0264

PROCEDURAL BLANK (PB):	A procedural blank (PB) was prepared with each analytical batch. Blanks were analyzed to ensure the sample extraction and analysis methods were free of contamination.
	07-0264 – No exceedences noted.
	Comments – None.
LABORATORY CONTROL SAMPLE (LCS):	A laboratory control sample (LCS) was prepared with each analytical batch. The percent recoveries of target analytes were calculated to measure data quality in terms of accuracy. 07-0264 – No exceedences noted.
	Comments – None.
SURROGATE RECOVERY:	Two surrogate compounds were added prior to extraction, including o-terphenyl and 5a- androstane. The recovery of the surrogate compound was calculated to measure data quality in terms of accuracy (extraction efficiency).
	07-0264 – No exceedences noted.
	Comments – None.
CONTROL OIL:	A control oil (North Slope Crude) was prepared with the analytical batch. The percent difference (PD) between the measured value and the target value was calculated to measure data quality in terms of accuracy.
	07-0264 – No exceedences noted.
	Comments – None.
CALIBRATIONS:	The GC/FID is calibrated with a minimum 5 level curve for all compounds. The percent
	relative standard deviation (% RSD) between RF for the individual target analytes must be $\leq 25\%$, and the mean RSD of all target analytes must be $\leq 20\%$. Each batch of samples analyzed is bracketed by a continuing calibration verification (CCV) sample, run at a frequency of minimally every 12 hours. The PD between the true value and the CCV should be $\leq 25\%$ for individual analytes. Additionally an initial calibration check (ICC) sample is run immediately after each initial calibration. The PD between the ICC and the initial calibration should be $\leq 25\%$.
	07-0264 – No exceedences noted.

Comments - None.

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Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	MW-9
Battelle ID	Q0540-P
Sample Type	
Collection Date	SA
Extraction Date	10/03/07
Analysis Date	10/18/07
Analytical Instrument	10/19/07
% Moisture	FID
% Lipid	NA
Matrix	NA
Sample Size	NAPL
Size Unit-Basis	53.70
Units	MG_OIL
o nue	UG/MG_OIL
n-Nonane	U
n-Decane	U
n-Undecane	U
n-Dodecane	U
n-Tridecane	U
soprenoid RRT 1380	U
n-Tetradecane	Ŭ
soprenoid RRT 1470	U
n-Pentadecane	U
1-Hexadecane	U
Norpristane (1650)	U
Hepladecane	U
Pristane	12.89
1-Octadecane	5.95
Phytane	8.62
-Nonadecane	U
Eicosane	Ú
-Heneicosane	Ŭ
-Docosane	Ũ
-Tricosane	Ŭ
-Tétracosane	ũ
-Pentacosane	ŭ
Hexacosane	Ŭ
-Heptacosane	Ŭ
Octacosane	Ŭ
Nonacosane	Ŭ
Triacontane	ŭ
Hentriacontane	Ŭ
Dotriacontane	Ŭ
Tritriacontane	Ŭ
Tetratriacontane	Ŭ
Pentatriacontane	U
Hexatriacontane	ŭ
Heptatriacontane	Ŭ
Octatriacontane	υ
Nonatriacontane	U
Tetracontane	
PH(total)	1025 45
, (lolo)	1035.45
urrogate Recoveries (%)	
Terphenyl -androstane	100 111

Battelle The Business of Innovation

Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	Procedural Blank	
Battelle ID		
	BL057PB-P	
Sample Type	PB	
Collection Date	10/18/07	
Extraction Date	10/18/07	
Analysis Date	10/19/07	
Analytical Instrument	FID	
% Moisture		
	NA	
% Lipid	NA	
Matrix	OIL	
Sample Size	50.00	
Size Unit-Basis	MG_OIL	
Units	UG/MG_OIL	
n-Nonane	U	
n-Decane	Ŭ	
n-Undecane	ŭ	
n-Dodecane	Ŭ	
n-Tridecane		
	U	
Isoprenoid RRT 1380	U	
n-Tetradecane	U	
Isoprenoid RRT 1470	U	
n-Pentadecane	U	
n-Hexadecane	U	
Norpristane (1650)	U	
n-Heptadecane	U	
Pristane	Ū	
n-Octadecane	ŭ	
Phytane	Ŭ	
n-Nonadecane	Ŭ	
n-Eicosane		
n-Heneicosane	U	
n-Docosane	U	
	U	
n-Tricosane	U	
n-Tetracosane	U	
n-Pentacosane	U	
n-Hexacosane	U	
n-Heptacosane	U	
n-Octacosane	U	
n-Nonacosane	U	
n-Triacontane	ŭ	
n-Hentriacontane	Ŭ	
n-Dotriacontane	Ŭ	
n-Tritriacontane		
n-Tetratriacontane	U	
n-Pentatriacontane	U	
	U	
n-Hexatriacontane	U	
n-Heptatriacontane	U	
n-Octatriacontane	U	
n-Nonatriacontane	U	
n-Tetracontane	U	
TPH(total)	U	
Surrogate Recoveries (%)		
O.T. I.I.I		
O-Terphenyl	98	
5a-androstane	103	
5a-androstane		

Battelle

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Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	Laboratory Control Sample					
Battelle ID	BL058LCS-P					
Sample Type	LCS					
Collection Date	10/18/07					
Extraction Date	10/18/07					
Analysis Date	10/19/07					
Analytical Instrument	FID					
% Moisture	NA					
% Lipid	NA					
Matrix	OIL					
Sample Size	NA					
Size Unit-Basis	NA					
Units	UG		Target	% Recovery	Qualifier	
n-Nonane	38.26		40.00			
n-Decane	38.3		40.00	96		
n-Undecane	37.71		40.00	96		
n-Dodecane	36.73		40.00	94		
n-Tridecane	36.4		40.00	92		
Isoprenoid RRT 1380		U	40.00	91		
n-Tetradecane	36.42		40.00	.04		
Isoprenoid RRT 1470		J	40.00	91		
n-Pentadecane	36.18	5	40.00	90		
n-Hexadecane	36.07		40.00	90		
Norpristane (1650)		J	40.00	90		
n-Heptadecane	36.22		40.00	91		
Pristane	36.94		40.00	92		
n-Octadecane	35.13		40.00	88		
Phytane	35.29		40.00	88		
n-Nonadecane	36.27		40.00	91		
n-Eicosane	36.3		40.00	91		
n-Heneicosane	35.77		40.00	89		
n-Docosane	36.9		40.00	92		
n-Tricosane	35.92		40.00	90		
n-Tetracosane	36.03		40.00	90		
n-Pentacosane	35.68		40.00	89		
n-Hexacosane	35.43		40.00	89		
n-Heptacosane	35.89		40.00	90		
n-Octacosane	34.39		40.00	86		
n-Nonacosane	35.09		40.00	88		
n-Triacontane	34.88		40.00	87		
n-Hentriacontane	34,33		40.00	86		
n-Dotriacontane	35.38		40.00	88		
n-Tritriacontane	34.61		40.00	87		
n-Tetratriacontane	34.9		40.00	87		
n-Pentatriacontane	36.01		40.00	90		
n-Hexatriacontane	33.65		40.00	84		
n-Heptatriacontane	28.76		40.00	72		
n-Octatriacontane n-Nonatriacontane	27.14		40.00	68		
n-Tetracontane	23.62		40.00	59		
TPH(total)	21.25		40.00	53		
(rentional)	U					

Surrogate Recoveries (%)

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O-Terphenyl	96
5a-androstane	100

Battelle

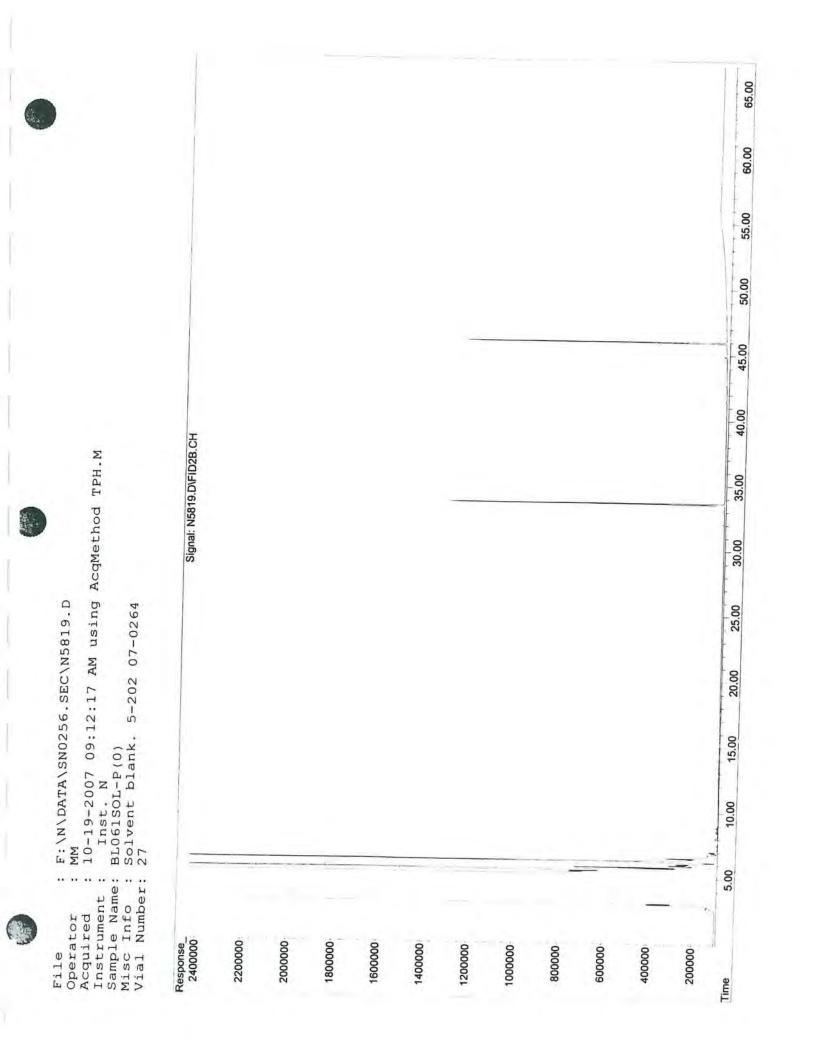
The Business of Innovation

Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Project Number: N007097-0001				
05-410	GO98: North Slope			
Client ID	Crude			
Battelle ID	BL059NSC-P			
Sample Type	NSC			
Collection Date	10/18/07			
Extraction Date	10/18/07			
Analysis Date	10/19/07			
Analytical Instrument	FID			
% Moisture	NA			
% Lipid	NA			
Matrix	OIL			
Sample Size	5.04			
Size Unit-Basis	MG OIL			
Units	UG/MG_OIL	Target %	Difference C	Qualifier
n-Nonane	1.05			
1-Decane	4.65 4.65	4.67	0.4	
n-Undecane	4.65	4.95	6.1	
1-Dodecane		4.51	2.1	
n-Tridecane	4.13	4.58	9.8	
soprenoid RRT 1380	4.13	4.19	1.4	
n-Tetradecane	0.96	0.96	0.2	
soprenoid RRT 1470	3.86	3.92	1.5	
Pentadecane	1.48	1.53	3.4	
-Hexadecane	3.98	3.99	0.3	
lorpristane (1650)	3.53	3.64	3.0	
-Heptadecane	1.12	1.14	1.9	
ristane	3.12	3.08	1.4	
-Octadecane	2.32	2.28	1.7	
hytane	2.96	2.80	5.8	
-Nonadecane	1,44	1.66	13.2	
-Eicosane	2.61	2.54	2.7	
-Heneicosane	2.58	2.50	3.1	
-Docosane	2.27	2.42	6.2	
Tricosane	2.31	2.25	2.6	
Tetracosane	2.05	2.05	0.0	
Pentacosane	1.95	1.95	0.1	
Hexacosane	1.78	1.80	0.9	
Heptacosane	1.64	1.64	0.0	
Octacosane	1.25	1.23	1.5	
Nonacosane	0.98	1.00	2.4	
Triacontane	0.79	0.87	9.4	
Hentriacontane	0.67	0.67	0.1	
Dotriacontane	0.64	0.61	5.5	
Tritriacontane		0.47	7.7	
Tetratriacontane	0.43	0.40	7.8	
Pentatriacontane	0.34	0.37	8.5	
Hexatriacontane	0.41	0.38	8.4	
Heptatriacontane	0.25 J	0.24	6.1	
Octatriacontane	0.21 J	0.21	0.0	
Nonatriacontane	0.2 J	0.21	2.8	
Tetracontane	0.15 J	0.15	2.5	
PH(total)	0.16 J	0.16	1.0	
i i i i i i i i i i i i i i i i i i i	478.35	578.97	17.4	

Surrogate Recoveries (%)

O-Terphenyl	97
5a-androstane	104





File : F:\N\DATA\SN0256.SEC\N5821.D Operator : MM Acquired : 19 Oct 2007 10:31 am using AcgMethod TPH.M Instrument : Inst. N Sample Name: BL062SOL-P(0) Misc Info : Solvent blank. 5-202 07-0264 Vial Number: 28

															t t t t t
				-											
Ð															
Signal: N5821.D\FID2B.CH											_				DE DO
Signal: N5															UU US
															25.00
															20.00
															15.00
															10.00
14(m)													=	1.1	5.00
Response160000015000001500000150000015000001500000150000015000001500000150000015000001500000001500000001500000000	1400000	1300000	1200000	1100000	100000	000006	800000	700000	600000	50000	400000	300000	200000	100000	Time



11:51 am using AcqMethod TPH.M 19 Oct 2007 11:51 am using Acc Inst. N BL057PB-P-FID(2) Procedural Blank 5-202 07-0264 29 F:\N\DATA\SN0256.SEC\N5823.D MM File : I Operator : N Acquired : A Instrument : Sample Name: E Misc Info : E Vial Number: 2

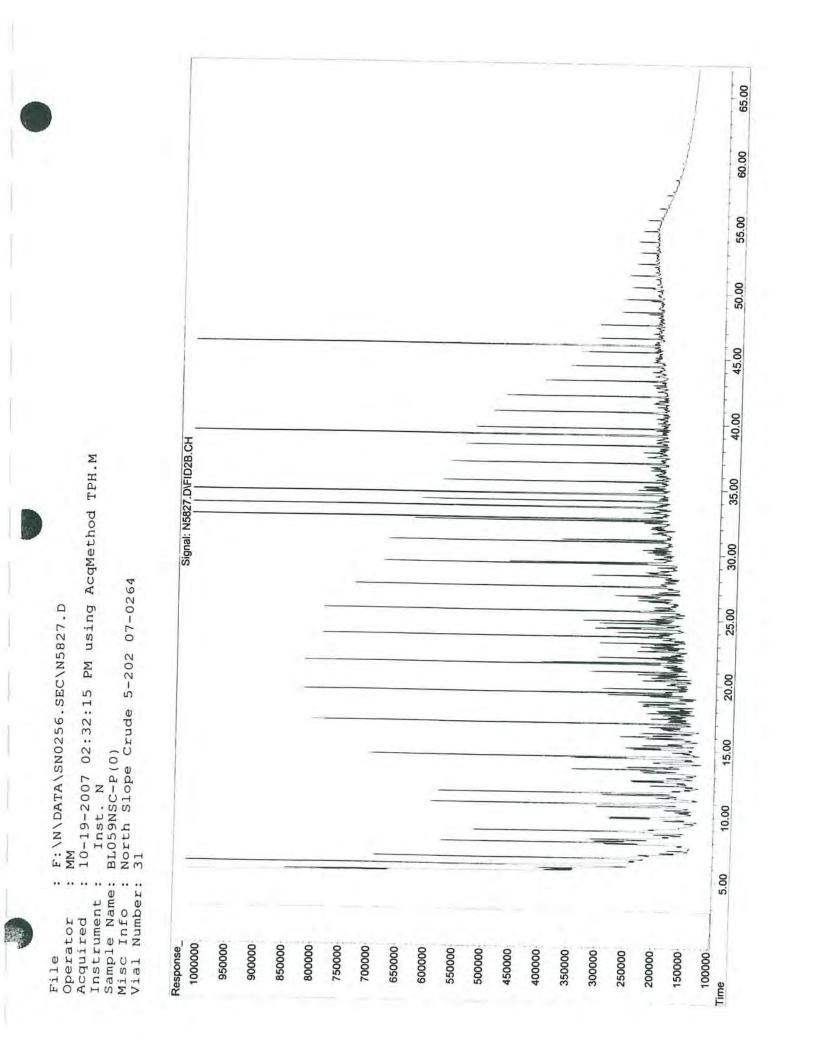
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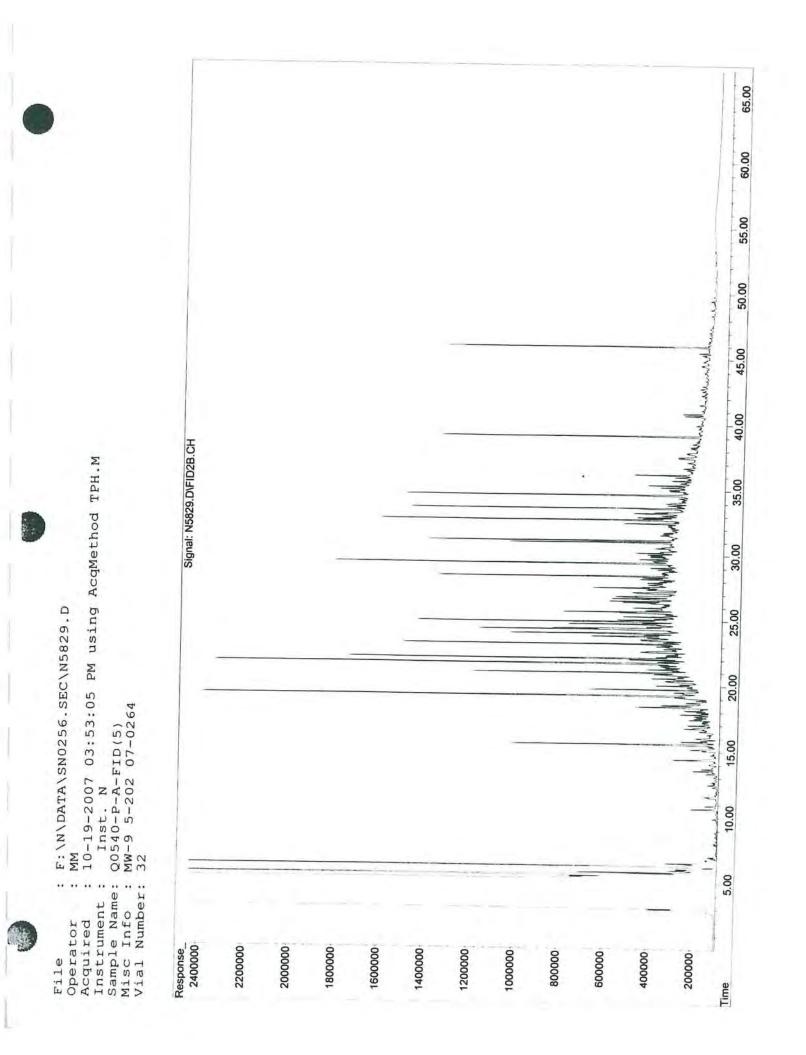


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SHC and TPH – SEDIMENT QA/QC SUMMARY Batch 07-0266

PROJECT PARAME LABORA' MATRIX: SAMPLE	TER: FORY:	Saturated H Battelle, Du Sediment Six soil sam Laboratory of taken and th temperature 2°) at 0.6°C. outside of th only on the i client was no analysis. Sa	er – Gas Works I ydrocarbons (SE xbury, MA ples and 1 NAP) on 10/9/07. Upo e samples were 1 of the cooler upo Also, it was no e container as w nside of the bub otified on 10/16/ mples were store aration could beg	IC) and Total P L were received on Receipt of the logged into the l on receipt was s ted that sample ell as the inner s ble wrap and di 07. The labora ed in an access-J	at Battelle Du samples, the t laboratory and lightly below t Q0540 (MW-9 side of the bubl d not touch any tory was instru	xbury Operatio emperature of t given unique II he acceptable r blad some oil ble wrap. The of the other sa cted to proceed	ns (BDO) the cooler was Ds. The ange $(4^{\circ}C \pm$ residue on the pil residue was mples. The with the
	Reference Method	Method Blank	Surrogate Recovery	LCS Recovery	MS Recovery	MS/MSD Precision	Control Oil % Diff
SHC and TPH	General NS&T	<5xMDL	40-120% Recovery	40-120% Recovery	40-120% Recovery	≤30% RPD	PD,30% for 90% of the

MS target spi	ke
must be >5x	
background	

analytes

METHOD:

Soil samples were extracted following general NS&T methods. Approximately 5-8 g of sample was spiked with SHC and PAH surrogates and serial extracted three times with dichloromethane using orbital shaker table techniques. The combined extracts were dried over anhydrous sodium sulfate and concentrated by Kuderna-Danish and nitrogen evaporation techniques. The sample extracts were split in half: one-half of the extract was removed for archiving; the other half was processed through an alumina gravity column to isolate the hydrocarbon fractions of interest. The weights of the resulting extracts were determined gravimetrically. The extracts were concentrated to 1 mL, split, and spiked with internal standard (IS). The pre-injection volume and/or extract split were adjusted to 5mg/mL. One extract was submitted for PAH and the second extract was submitted for SHC and TPH analysis.

SHC and TPH were measured by gas chromatography with flame ionization detection (GC/FID). An initial calibration consisting of target analytes was completed prior to analysis to demonstrate the linear range of the analysis. Calibration verification was performed at the beginning and end of each 12 hour period in which samples were analyzed. Concentrations of SHC and TPH were calculated by the internal standard method. Normal alkanes were quantified using the average RF generated from the initial calibration. TPH concentrations were quantified suing the average RF of nC8 through nC40. Isoprenoid hydrocarbon concentrations were quantified using the average RF of nc8 through nC40. Isoprenoid hydrocarbon concentrations were quantified using the average RF of nc8 through nC40. Isoprenoid hydrocarbon concentrations were quantified using the average RF of nc8 through nC40. Isoprenoid hydrocarbon concentrations were quantified using the average RF of nc8 through nC40. Isoprenoid

HOLDING TIMES: Samples were prepared for analysis in three analytical batches and were extracted within 30 days of sample collection analyzed within 40 days of extraction. The samples from this batch were re-extracted from 07-0259 because of poor surrogate recoveries.

Batch	Extraction Date	Analysis Date
07-0266	10/23/2007	10/30/2007 - 10/31/2007

SHC and TPH – SEDIMENT QA/QC SUMMARY Batch 07-0266

PROCEDURAL BLANK (PB):	A procedural blank (PB) was prepared with each analytical batch. Blanks were analyzed to ensure the sample extraction and analysis methods were free of contamination.
	07-0266 – No exceedences noted.
	Comments – None.
LABORATORY CONTROL SAMPLE (LCS):	A laboratory control sample (LCS) was prepared with each analytical batch. The percent recoveries of target analytes were calculated to measure data quality in terms of accuracy.
	07-0266 – No exceedences noted.
	Comments – None.
SURROGATE RECOVERY:	Two surrogate compounds were added prior to extraction, including o-terphenyl and 5a- androstane. The recovery of the surrogate compound was calculated to measure data quality in terms of accuracy (extraction efficiency).
	07-0266 – Three of the four samples extracted in this batch failed SIS recovery criteria with high recoveries.
	Comments – These samples have been extracted twice with similar results. The issue appears to be matrix related rather than extraction related as the PB and LCS (samples without matrix) did not exhibit similar trends.
CONTROL OIL:	A control oil (North Slope Crude) was prepared with the analytical batch. The percent difference (PD) between the measured value and the target value was calculated to measure data quality in terms of accuracy.
	07-0266 – nC13 had a PD of 48%, but was "ME" qualified indicating that an interferent was identified. The control oil still passed the MQO of the PD being \leq 30% for 90% of the analytes.
	Comments – None.
CALIBRATIONS:	The GC/FID is calibrated with a minimum 5 level curve for all compounds. The percent
	relative standard deviation (% RSD) between RF for the individual target analytes must be $\leq 25\%$, and the mean RSD of all target analytes must be $\leq 20\%$. Each batch of samples analyzed is bracketed by a continuing calibration verification (CCV) sample, run at a frequency of minimally every 12 hours. The PD between the true value and the CCV should be $\leq 25\%$ for individual analytes. Additionally an initial calibration check (ICC) sample is run immediately after each initial calibration. The PD between the ICC and the initial calibration should be $\leq 25\%$.

07-0266 - No exceedences noted.

Comments - None.

Battelle

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Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	SB-8 S5 9-10.5	SB-13 2.5-4.0	SB-12A S3 5-6.5	GP-9 7-8
Battelle ID	Q0542-P1	Q0543-P1	00544.04	Section 1
Sample Type	SA		Q0544-P1	Q0546-P1
Collection Date		SA	SA	SA
	09/18/07	09/20/07	09/20/07	09/18/07
Extraction Date	10/23/07	10/23/07	10/23/07	10/23/07
Analysis Date	10/30/07	10/31/07	10/31/07	10/31/07
Analytical Instrument	FID	FID	FID	FID
% Moisture	8.06	16.77	13.98	
% Lipid	NA	NA		49.24
Matrix	SOIL		NA	NA
Sample Size		SOIL	SOIL	SOIL
Size Unit-Basis	4.78	4.45	4.63	2.84
Units	G_DRY	G_DRY	G_DRY	G_DRY
onto	UG/G_DRY	UG/G_DRY	UG/G_DRY	UG/G_DRY
n-Nonane	U	U	U	U
n-Decane	U	Ŭ	ŭ	
n-Undecane	Ū	Ŭ		U
n-Dodecane	Ŭ	υ	<u>U</u> -	U
n-Tridecane	ŭ		U	U
Isoprenoid RRT 1380		U	U	U
n-Tetradecane	U	U	U	U
Isoprenoid RRT 1470	U	U	U	U
	U	U	U	U
n-Pentadecane	U	U	U	U
n-Hexadecane	U	U	U	U
Norpristane (1650)	28.63	U	Ú	Ŭ
n-Heptadecane	U	U	U	Ŭ
Pristane	41.71	U	Ŭ	Ŭ
n-Octadecane	U	Ŭ	Ŭ	Ŭ
Phytane	25.27	Ū	ũ	Ŭ
n-Nonadecane	U	Ũ	Ŭ	
n-Eicosane	Ŭ	ŭ	U U	U
n-Heneicosane	Ŭ	Ŭ		U
n-Docosane	Ŭ	U	U	U
n-Tricosane	ŭ		u	U
n-Tetracosane	Ŭ	U	U	U
n-Pentacosane	U	U	U	U
n-Hexacosane		U	U	U
n-Heptacosane	U	U	U	υ
n-Octacosane	U	υ	U	U
n-Nonacosane	U	U	U	U
	u	U	U	U
n-Triacontane	U	U	U	Ŭ
n-Hentriacontane	U	U	U	ũ
n-Dotriacontane	U	U	U	Ŭ
n-Tritriacontane	U	U	U	Ŭ
n-Tetratriacontane	U	U	Ŭ	Ŭ
n-Pentatriacontane	U	Ŭ	ŭ	U
n-Hexatriacontane	U	ũ	Ŭ	
n-Heptatriacontane	U	ŭ	Ŭ	0
n-Octatriacontane	υ	Ŭ	ŭ	U
n-Nonatriacontane	Ũ	11	71	U
n-Tetracontane	Ŭ	Ŭ	U	U
TPH(total)	4533.08	10664.19	U 45954.94	U 53535.09
Surrogate Recoveries (%)				
O-Terphenyl	150.11			
5a-androstane	150 N	63	165 N	148 N
	142 N	52	150 N	152 N

Battelle

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Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	Procedural Blank	
Battelle ID	BL065PB-P	
Sample Type		
Collection Date	PB	
Extraction Date	10/23/07	
	10/23/07	
Analysis Date	10/30/07	
Analytical Instrument	FID	
% Moisture	22.01	
% Lipid	NA	
Matrix	SEDIMENT	
Sample Size	4.20	
Size Unit-Basis	G DRY	
Units	UG/G_DRY	
n-Nonane	U	
n-Decane	Ŭ	
n-Undecane	Ŭ	
n-Dodecane	U U	
n-Tridecane	U	
Isoprenoid RRT 1380		
n-Tetradecane	U	
Isoprenoid RRT 1470	U	
n-Pentadecane	U	
	U	
n-Hexadecane	U	
Norpristane (1650)	U	
n-Heptadecane	U	
Pristane	U	
n-Octadecane	U	
Phytane	U	
n-Nonadecane	U	
n-Eicosane	U	
n-Heneicosane	U	
n-Docosane	Ŭ	
n-Tricosane	U	
n-Tetracosane	U	
n-Pentacosane	Ŭ	
n-Hexacosane	Ŭ	
n-Heptacosane	Ŭ	
1-Octacosane	Ŭ	
I-Nonacosane	Ŭ	
-Triacontane	Ű	
-Hentriacontane	U	
Dotriacontane	UUU	
-Tritriacontane		
Tetratriacontane	U.	
-Pentatriacontane	U U	
-Hexatriacontane		
-Heptatriacontane	U	
-Octatriacontane	- U	
-Nonatriacontane	U	
	U	
-Tetracontane	υ	
PH(total)	U	
urrogate Recoveries (%)		

O-Terphenyl 5a-androstane

106 102

Battelle The Business of Innovation

Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	060208-03: Sand, White Quartz, -50+70					
Battelle ID	BL066LCS-P					
Sample Type	LCS					
Collection Date	10/23/07					
Extraction Date	10/23/07					
Analysis Date						
Analytical Instrument	10/30/07 FID					
% Moisture	NA					
% Lipid	NA					
Matrix	SEDIMENT					
Sample Size	SEDIMENT					
Size Unit-Basis	NA					
Units	UG		Target 9	& Recovery	Qualifier	
All and					Gouinor	
n-Nonane	101.24		100.00	101		
n-Decane	101.13		100.00	101		
n-Undecane	102.74		100.00	103		
n-Dodecane	101.86		100.00	102		
n-Tridecane	105.12	1	100.00	105		
soprenoid RRT 1380 n-Tetradecane		U	100.000			
	101.67		100.00	102		
soprenoid RRT 1470 n-Pentadecane	123.5	U				
-Hexadecane	100.17		100.00	100		
Vorpristane (1650)	100.23		100.00	100		
n-Heptadecane		U				
Pristane	101.57		100.00	102		
-Octadecane	101.94		100.00	102		
Phytane	100.88		100.00	101		
-Nonadecane	98.48 102.23		100.00	98		
-Eicosane	102.23		100.00	102		
-Heneicosane	100.28		100.00	102		
-Docosane	102.83		100.00	100 103		
-Tricosane	99.87		100.00	100		
Tetracosane	100.28		100.00	100		
-Pentacosane	99.02		100.00	99		
-Hexacosane	99.28		100.00	99		
-Heptacosane	100.24		100.00	100		
Octacosane	101.45		100.00	101		
Nonacosane	98		100.00	98		
Triacontane	97.65		100.00	98		
Hentriacontane	95.49		100.00	95		
Dotriacontane	96.81		100.00	97		
Tritriacontane	93.15		100.00	93		
Tetratriacontane	91.62		100.00	92		
Pentatriacontane	93.43		100.00	93		
Hexatriacontane	86.24		100.00	86		
Heptatriacontane	84		100.00	84		
Octatriacontane	84.54		100.00	85		
Nonatriacontane	77.68		100.00	78		
Tetracontane	74.91		100.00	75		
PH(total)		U				

Surrogate Recoveries (%)

O-Terphenyl	118
5a-androstane	119

Battelle The Business of Innovation

Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	GO98: North Slope Crude				
	Crude				
Battelle ID	BL087NSC-P				
Sample Type	NSC				
Collection Date	10/30/07				
Extraction Date	10/30/07				
Analysis Date	10/30/07				
Analytical Instrument	FID				
% Moisture	NA				
% Lipid	NA				
Matrix	OIL				
Sample Size	5.04				
Size Unit-Basis	MG OIL				
Units	UG/MG_OIL	Target %	Difference	Qualifier	
Nonane					
-Decane	4.93	4.67	5.6		
Undecane	4.62	4.95	6.7		
n-Dodecane	4.48	4.51	0.6		
n-Tridecane	4.54	4.58	0.8		
soprenoid RRT 1380	6.2 ME	4.19	48.0	N	
Tetradecane	0.92	0.96	4.3		
soprenoid RRT 1470	4.23	3.92	7.9		
-Pentadecane	1.59	1.53	3.7		
-Hexadecane	3.88	3.99	2.8		
lorpristane (1650)	3.76 1.08	3.64	3.3		
-Heptadecane	3.33	1.14	5.4		
ristane	2.28	3.08	8.2		
-Octadecane	2.28	2.28	0.0		
hytane	1.52	1.66	1.3		
-Nonadecane	2,7	2.54	8.4 6.3		
-Eicosane	2.43	2.54	2.9		
Heneicosane	2.35	2.42	2.9		
Docosane	2.29	2.25	1.7		
Tricosane	2.01	2.05	2.0		
Tetracosane	1.96	1.95	0.6		
Pentacosane	1.65	1.80	8.1		
Hexacosane	1.53	1.64	6.7		
Heptacosane	1.24	1.23	0.7		
Octacosane	1.01	1.00	0.6		
Nonacosane	0.84	0.87	3.7		
Triacontane	0.67	0.67	0.1		
Hentriacontane	0.59	0.61	2.8		
Dotriacontane	0.49	0.47	5.2		
Tritriacontane Tetratriacontane	0.38	0.40	4.8		
Pentatriacontane	0.34	0.37	8.5		
Hexatriacontane	0.35	0.38	7.4		
Heptatriacontane	0.24 J	0.24	1.8		
Dctatriacontane	0.21 J	0.21	0.0		
Nonatriacontane	0.23 J	0.21	11.8		
fetracontane	0.17 J	0.15	10.4		
'H(total)	0.18 J	0.16	11.4		
. dramad	495.56	578.97	14.4		

Surrogate Recoveries (%)

O-Terphenyl	114
5a-androstane	110

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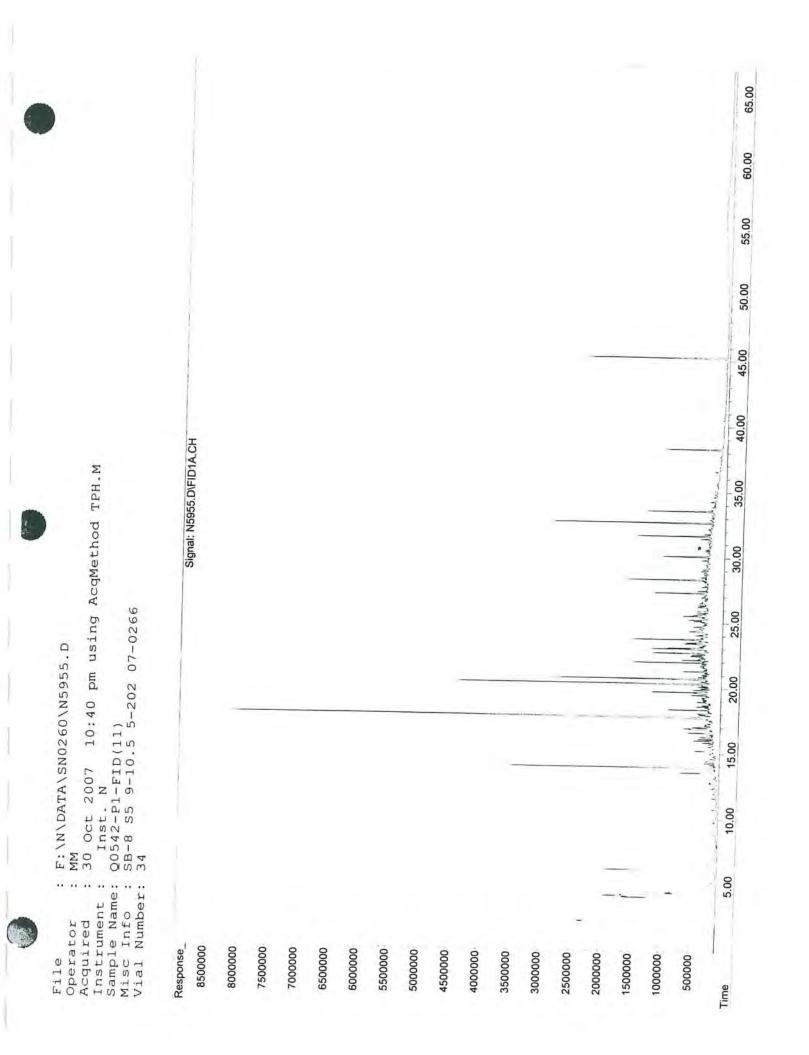
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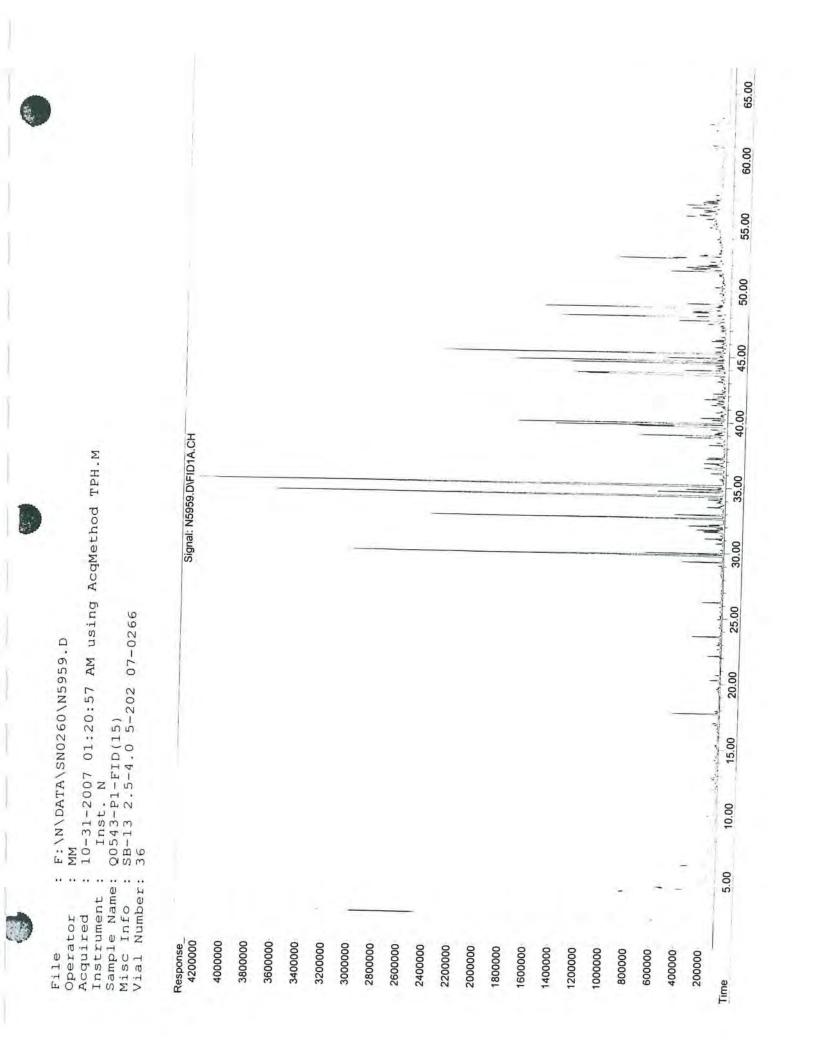
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	F:\N\DATA\SN0260\N5953.D MM	10-30-2007 Tret M	87NSC-F	North Slope Crude 5 33										-			10.00
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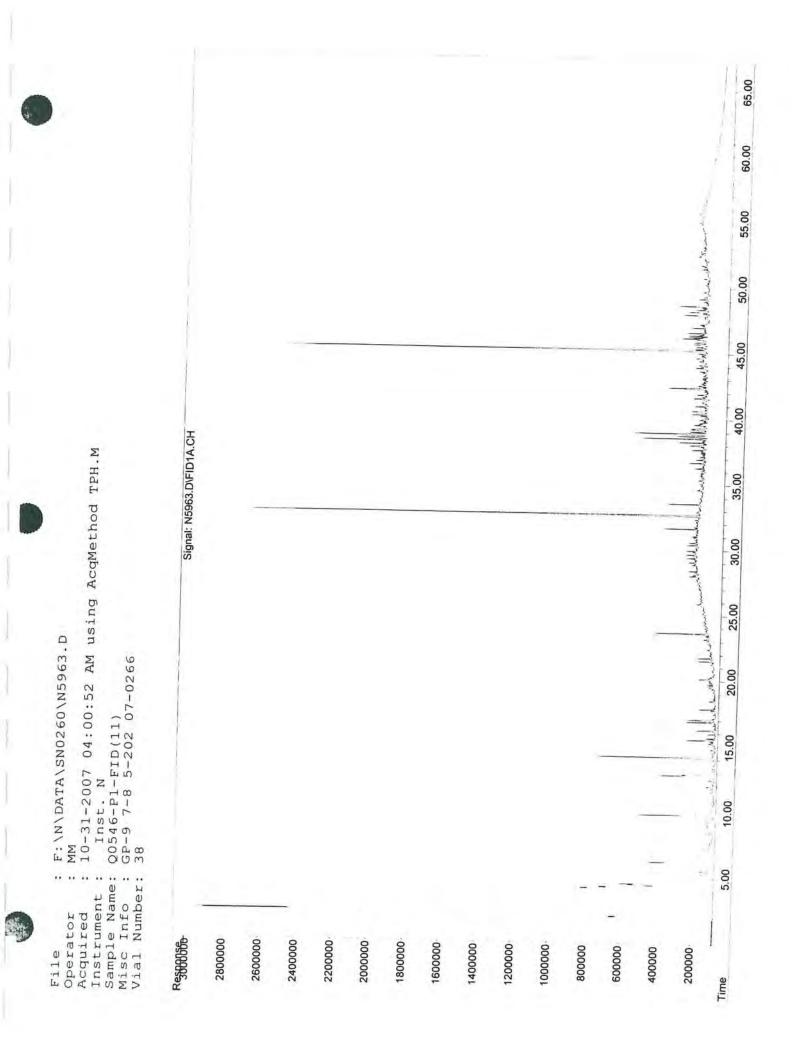
MHT	
F:\N\DATA\SN0260\N5961.D MM 10-31-2007 02:40:32 AM using AcaMethod TPH.M	
1.D M using	07-0266
F:\N\DATA\SN0260\N5961 MM 10-31-2007 02:40:32 AM	1 -FID(11) 8 5-6.5 5-202 07-0266
ATA\SN02 2007 02	Inst. N Q0544-P1-FID(J SB-12A S3 5-6. 37
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Response	9500000	8000006	850000	8000000	750000	700000	650000	8000000	550000	500000	4500000	4000000	350000	3000000	250000	2000000	1500000	1000000	

65.00

60.00

55.00



PAH Data and Histograms Biomarker Data and EICPs

PAH and Biomarker – SEDIMENT QA/QC SUMMARY Batch 07-0259

PROJECT PARAME LABORA' MATRIX: SAMPLE	TER: FORY:	Polycyclic A Battelle, Du: Sediment Six soil sam Laboratory of taken and the temperature 2°) at 0.6°C. outside of th only on the i client was no analysis. Sam	xbury, MA ples and 1 NAPI on 10/9/07. Upo e samples were 1 of the cooler up Also, it was no e container as w nside of the bub otified on 10/16/	L were received n Receipt of th ogged into the on receipt was ted that sample ell as the inner ble wrap and d 07. The labor ed in an access-	and Biomarkers d at Battelle Duy te samples, the te laboratory and g slightly below th e Q0540 (MW-9 side of the bubb id not touch any atory was instruc- limited walk-in	abury Operation emperature of t given unique II ne acceptable r) had some oil ole wrap. The o of the other sa cted to proceed	he cooler was Ds. The ange $(4^{\circ}C \pm$ residue on the bil residue was mples. The with the
	Reference Method	Method Blank	Surrogate Recovery	LCS Recovery	MS Recovery	MS/MSD Precision	Control Oil % Diff
SHC and TPH	General NS&T	<5xMDL	40-120% Recovery	40-120% Recovery	40-120% Recovery MS target spike	≤30% RPD	PD,30% for 90% of the analytes

METHOD:

HOLDING TIMES: Soil samples were extracted following general NS&T methods. Approximately 5-8 g of sample was spiked with SHC, PAH, and biomarker surrogates and serial extracted three times with dichloromethane using orbital shaker table techniques. The combined extracts were dried over anhydrous sodium sulfate and concentrated by Kuderna-Danish and nitrogen evaporation techniques. The sample extracts were split in half: one-half of the extract was removed for archiving; the other half was processed through an alumina gravity column to isolate the hydrocarbon fractions of interest. The weights of the resulting extracts were determined gravimetrically. The extracts were concentrated to 1 mL, split, and spiked with internal standard (IS). The pre-injection volume and/or extract split were adjusted to 5mg/mL. One extract was submitted for PAH and the second extract was submitted for SHC and TPH analysis.

PAH and petroleum biomarkers were measured by gas chromatography -mass spectrometry (GC/MS) in the selected ion mode (SIM). An initial calibration consisting of target analytes was completed prior to analysis to demonstrate the linear range of the analysis. Calibration verification was performed at the beginning and end of each 12 hour period in which samples were analyzed. Concentrations of the PAH and petroleum biomarkers were calculated by the internal standard method. Target PAH were quantified using the average RF generated from the initial calibration. The alkyl homologue PAH series were assigned the RF of the parent PAH, Steranes were assigned the RF of cholestane, and triterpanes were assigned the RF of Moretane.

Note: the reporting limit for the alkyl benzene compounds is orders of magnitude higher than the reporting limits for the rest of the PAH compounds.

Samples were prepared for analysis in three analytical batches and were extracted within 30 days of sample collection analyzed within 40 days of extraction.

 Batch
 Extraction Date
 Analysis Date

 07-0259
 10/15/2007
 10/24/2007

PAH and Biomarker – SEDIMENT QA/QC SUMMARY Batch 07-0259

PROCEDURAL BLANK (PB):	A procedural blank (PB) was prepared with each analytical batch. Blanks were analyzed to ensure the sample extraction and analysis methods were free of contamination.
	07-0259 – No exceedences noted.
	Comments – None.
LABORATORY CONTROL SAMPLE (LCS):	A laboratory control sample (LCS) was prepared with each analytical batch. The percent recoveries of target analytes were calculated to measure data quality in terms of accuracy.
	07-0259 – No exceedences noted.
	Comments – None.
SURROGATE RECOVERY:	Five surrogate compounds were added prior to extraction, including naphthalene-d8, acenaphthene-d10, phenanthrene-d10, benzo(a)pyrene-d12, and 5b(H)-chloane. The recovery of the surrogate compound was calculated to measure data quality in terms of accuracy (extraction efficiency).
	07-0259 – Several surrogates are over-recovered in the samples. Benzo(a)pyrene-d12 is out in the PB and the NSC, though the NSC still passes for all the analytes and this does not affect the authentic samples.
	Comments – The exceedences in the authentic samples is most likely attributed to the high levels of target analytes found in the samples and should have not affect on the values reported for the target analytes.
CONTROL OIL:	A control oil (North Slope Crude) was prepared with the analytical batch. The percent difference (PD) between the measured value and the target value was calculated to measure data quality in terms of accuracy.
	07-0259 – No exceedences noted.
	Comments – None.
CALIBRATIONS:	The GC/MS is calibrated with a minimum 5 level curve for all compounds. The percent relative standard deviation (% RSD) between RF for the individual target analytes must be $\leq 25\%$, and the mean RSD of all target analytes must be $\leq 15\%$. Each batch of samples analyzed is bracketed by a continuing calibration verification (CCV) sample, run at a frequency of minimally every 12 hours. The PD between the true value and the CCV should be $\leq 25\%$ for individual analytes. Additionally an initial calibration check (ICC) sample is run immediately after each initial calibration. The PD between the ICC and the initial calibration should be $\leq 25\%$.

07-0259 - No exceedences noted.

Comments - None.

The Business of Innovation

Client ID	SB-2 S5 8-9.5	GP-12 S2 23-24
Battelle ID	Q0541-P	Q0545-P
Sample Type	SA	SA SA
Collection Date	09/17/07	09/20/07
Extraction Date	10/15/07	10/15/07
Analysis Date	10/24/07	10/24/07
Analytical Instrument	MS	
% Moisture	18.05	MS
% Lipid	NA	26.35
Matrix	SOIL	NA
Sample Size	4.59	SOIL
Size Unit-Basis	G DRY	3.93
Units	NG/G_DRY	G_DRY NG/G_DRY
C3-Alkylbenzenes	10085.26	5127.04
C4-Alkylbenzenes	15434.72	
C5-Alkylbenzenes	10512.82	3892.29
C6-Alkylbenzenes		742.61 J
Benzo(b)thiophene	12276.98	339 J
C1-benzo(b)thiophenes	9878.53	20344.59
C2-benzo(b)thiophenes	11507.34	7546.34
C3-benzo(b)thiophenes	14450.56	4669.98
C4-benzo(b)thiophenes	14482.43	1168.82
Naphthalene	8555.49	159.28
C1-Naphthalenes	280712.07 D	401296.71 D
C2-Naphthalenes	161435.32 D	106483 D
C3-Naphthalenes	200528.1 D	32509.83 D
C4-Naphthalenes	150550.6 D	6892.79 D
	78992.16 D	1297.11 D
C1-Biphenyls + Dibenzofuran	28932.64	16556.81
Biphenyl	8957.51	12549.72
C2-Biphenyls + C1-Dibenzofurans	45019.06	6521.65
Acenaphthylene	5745.65	15068.74
Acenaphthene	122429.23 D	6762.95
Dibenzofuran	7893.62	12352.38
Fluorene	64333.84 D	20199.05
C1-Fluorenes	37349.9 D	3928.84
C2-Fluorenes	32374.2 D	1332.36
C3-Fluorenes	20462.87 D	565.53
C4-Fluorenes	9721.05 D	233.13
Anthracene	54254.5 D	15041.8 D
Phenanthrene	187892.05 D	60533.18 D
C1-Phenanthrenes/Anthracenes	109645.54 D	12567.33 D
C2-Phenanthrenes/Anthracenes	55169.03 D	5974.74 D
C3-Phenanthrenes/Anthracenes	20750.58 D	1092.05 D
C4-Phenanthrenes/Anthracenes	5696.83 D	442.09
Retene	2259.18	891.41
Dibenzothiophene	15598.08	3943.26
C1-Dibenzothiophenes	18802.31	1433.37
C2-Dibenzothiophenes	16145.18	660.06
C3-Dibenzothiophenes	8872.71	234.5
C4-Dibenzothiophenes	3335.64	67.4
luoranthene	59196.4 D	22336.82
Pyrene	97480.76 D	20773.98
1-Fluoranthenes/Pyrenes	65145.99 D	12195.01
2-Fluoranthenes/Pyrenes	14924.46 D	3270.47
3-Fluoranthenes/Pyrenes	4915.67 D	
4-Fluoranthenes/Pyrenes	2180.92 D	988.68
CO-Benzo(b)naphthothiophenes	3776.25	465.25
1-Benzo(b)naphthothiophenes	4725.17	913.61
2-Benzo(b)naphthothiophenes	the Base of the State of the St	694.13
3-Benzo(b)naphthothiophenes	2125.13	221.74
4-Benzo(b)naphthothiophenes	1212.17	154.67
enzo(a)anthracene	536.8	36.21
hrysene	24074,78	9590.03
	22810.15	9771.77

The Business of Innovation

Client ID	SB-2 S5 8-9.5	GP-12 S2 23-24
Battelle ID	Q0541-P	00545 0
Sample Type	SA SA	Q0545-P
Collection Date	09/17/07	SA
Extraction Date	10/15/07	09/20/07
Analysis Date	10/15/07	10/15/07
Analytical Instrument		10/24/07
% Moisture	MS	MS
% Lipid	18.05	26.35
Matrix	NA	NA
Sample Size	SOIL	SOIL
Size Unit-Basis	4.59	3.93
Units	G_DRY	G_DRY
Units	NG/G_DRY	NG/G_DRY
C1-Chrysenes	16589.65	4494.91
C2-Chrysenes	5241.89	1459.72
C3-Chrysenes	2403.14	619.55
C4-Chrysenes	1114.53	241.79
Benzo(b)fluoranthene	12560.8	7467.49
Benzo(k)fluoranthene	17864.18	8774.71
Benzo(e)pyrene	15474.59	6186
Benzo(a)pyrene	28503.51	11388.72
Pervlene	4224.91	3288.78
Indeno(1,2,3-cd)pyrene	12042.45 D	5028.52 D
Dibenz(a,h)anthracene	2886.08 D	1140.83 D
Benzo(g,h,i)perylene	13776.39	7060.88
Total PAH	2089157.85	859307.45
		000007.40
Surrogate Recoveries (%)		
Naphthalene-d8	123 N	103
cenaphthene-d10	189 N	110
Phenanthrene-d10	94	
Phenanthrene-d10 Benzo(a)pyrene-d12	94 107 D	93 72 D

Battelle The Business of Innovation

Client ID	GO98: North Slope Crude			
Battelle ID	BL042NSC-P			
Sample Type	NSC			
Collection Date	10/17/07			
Extraction Date	10/17/07			
Analysis Date	10/24/07			
Analytical Instrument	MS			
% Moisture	NA			
% Lipid	NA			
Matrix	OIL			
Sample Size	5.04			
Size Unit-Basis	MG_OIL			
Units	MG/KG_OIL	Target %	Difference	Qualifier
C3-Alkylbenzenes	1616.97			
C4-Alkylbenzenes	1276.68			
C5-Alkylbenzenes	721.47			
C6-Alkylbenzenes	454.78			
Benzo(b)thiophene	12.64			
C1-benzo(b)thiophenes	42.96			
C2-benzo(b)thiophenes	76.93	95.74	19.6	
C3-benzo(b)thiophenes	99.29	132.67	25.2	
C4-benzo(b)thiophenes	96.03	96.72	0.7	
Naphthalene C1 Naphthalene	705.29	740.29	4.7	
C1-Naphthalenes	1418.33	1516.04	6.4	
C2-Naphthalenes C3-Naphthalenes	1865.4	2000.10	6.7	
C4-Naphthalenes	1459.07	1526.96	4.4	
C1-Biphenyls + Dibenzofuran	792.31 338.3	898.03	11.8	
Biphenyl	211	220.82	4.4	
C2-Biphenyls + C1-Dibenzofurans	502.68	220.02	4.4	
Acenaphthylene	002.00 U			
Acenaphthene	13.59	14.50	6.3	
Dibenzofuran	67.6	77.75	13.1	
Fluorene	98.21	92.51	6.2	
C1-Fluorenes	209.14	227.01	7.9	
C2-Fluorenes	298.65	367.09	18.6	
C3-Fluorenes	274.46	326.32	15.9	
C4-Fluorenes	179.5			
Anthracene	U			
Phenanthrene	237.55	249.49	4.8	
C1-Phenanthrenes/Anthracenes	507.25	549.17	7.6	
C2-Phenanthrenes/Anthracenes	596.02	642.72	7.3	
C3-Phenanthrenes/Anthracenes	423.45	446,11	5.1	
C4-Phenanthrenes/Anthracenes Retene	150.11	180.02	16.6	
Dibenzothiophene	43.19	10.000		
C1-Dibenzothiophenes	197.91	210.35	5.9	
C2-Dibenzothiophenes	376.95	409.03	7.8	
C3-Dibenzothiophenes	520.69	551.46	5.6	
C4-Dibenzothiophenes	420.01 232.92	471.36	10.9	
Fluoranthene	3.51	243.11	4.2	
Pyrene	11.92	12.99	8.2	
C1-Fluoranthenes/Pyrenes	69.22	70.92	2.4	
C2-Fluoranthenes/Pyrenes	128.43	117.89	8.9	
C3-Fluoranthenes/Pyrenes	146.1	137.25	6.4	
C4-Fluoranthenes/Pyrenes	111.01	101,60	0.4	
C0-Benzo(b)naphthothiophenes	40.92			
C1-Benzo(b)naphtholhiophenes	160.58			
C2-Benzo(b)naphthothiophenes	194.47			
C3-Benzo(b)naphthothiophenes	143.86			
C4-Benzo(b)naphthothiophenes	96.41			
Benzo(a)anthracene	4.52			

Battelle The Business of Innovation

Project Number: N007097-0001					
	GO98: North Slope				
Client ID	Crude				
Battelle ID	BL042NSC-P				
Sample Type	NSC				
Collection Date	10/17/07				
Extraction Date	10/17/07				
Analysis Date	10/24/07				
Analytical Instrument	MS				
% Moisture	NA				
% Lipid	NA				
Matrix	OIL				
Sample Size	5.04				
Size Unit-Basis	MG_OIL				
Units	MG/KG_OIL		Target % D	Difference	Qualifier
Chrysene	10.00		17.10		
C1-Chrysenes	48.26		47.18	2.3	
C2-Chrysenes	83.68 108.82		78.82	6.2	
C3-Chrysenes	108.82		102.67	6.0	
C4-Chrysenes	98.2 60.53		85.36	15.0	
Benzo(b)fluoranthene	6.48		61.99	2.4	
Benzo(k)fluoranthene	0.68	J	6.08	6.6	
Benzo(e)pyrene	14.65	J	10.00	10.7	
Benzo(a)pyrene	14.05	υ	12.88	13.7	
Perylene		U			
ndeno(1.2.3-cd)pyrene		ŭ			
Dibenz(a,h)anthracene	1.26	0			
Benzo(g,h,i)perylene	3.46		3.44	0.0	
Total PAH	11865.63		3.44	0.6	
	11003.03				
Surrogate Recoveries (%)					
lashingland 10					
Naphthalene-d8	113				
Acenaphthene-d10 Phenanthrene-d10	98				
	96				
Benzo(a)pyrene-d12	139	N			
5b(H)-Cholane					

The Business of Innovation

Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	Procedural Blank	
Battelle ID	DI 033DC D	
Sample Type	BL033PB-P	
	PB	
Collection Date	10/15/07	
Extraction Date	10/15/07	
Analysis Date	10/24/07	
Analytical Instrument	MS	
% Moisture	22.07	
% Lipid	NA	
Matrix	SEDIMENT	
Sample Size	11.03	
Size Unit-Basis		
Units	G_DRY NG/G DRY	
C3-Alkylbenzenes	U	
C4-Alkylbenzenes	U	
C5-Alkylbenzenes	U	
C6-Alkylbenzenes	U	
Benzo(b)thiophene	Ŭ	
C1-benzo(b)thiophenes	ŭ	
C2-benzo(b)thiophenes	Ŭ	
C3-benzo(b)thiophenes		
C4-benzo(b)thiophenes	u	
	U	
Naphthalene	0.82 J	
C1-Naphthalenes	0.12 J	
C2-Naphthalenes	U	
C3-Naphthalenes	U	
C4-Naphthalenes	U	
C1-Biphenyls + Dibenzofuran	Ŭ	
Biphenyl	Ŭ	
C2-Biphenyls + C1-Dibenzofurans	Ŭ	
Acenaphthylene	Ŭ	
Acenaphthene		
Dibenzofuran	U	
	U	
Fluorene	u	
C1-Fluorenes	U	
C2-Fluorenes	U	
C3-Fluorenes	U	
C4-Fluorenes	Ũ	
Anthracene	ŭ	
Phenanthrene	Ŭ	
C1-Phenanthrenes/Anthracenes	Ŭ	
C2-Phenanthrenes/Anthracenes		
C3-Phenanthrenes/Anthracenes	U	
C4-Phenanthrenes/Anthracenes	U	
Retene	U	
	U	
Dibenzothiophene	U	
C1-Dibenzothiophenes	U	
C2-Dibenzothiophenes	U	
C3-Dibenzothiophenes	Ū	
C4-Dibenzothiophenes	ŭ	
luoranthene	Ŭ	
Pyrene		
C1-Fluoranthenes/Pyrenes	U	
22-Fluoranthenes/Pyrenes	U	
	U	
3-Fluoranthenes/Pyrenes	U	
24-Fluoranthenes/Pyrenes	U	
0-Benzo(b)naphthothiophenes	U U	
1-Benzo(b)naphthothiophenes	U	
2-Benzo(b)naphthothiophenes	Ŭ	
3-Benzo(b)naphthothiophenes	ũ	
4-Benzo(b)naphthothiophenes	Ŭ	
	U.	
enzo(a)anthracene	11	
enzo(a)anthracene hrysene	U U	

The Business of Innovation

Client ID	Procedural Blank	
Battelle ID	BL033PB-P	
Sample Type	PB	
Collection Date	10/15/07	
Extraction Date	10/15/07	
Analysis Date	10/24/07	
Analytical Instrument	MS	
% Moisture	22.07	
% Lipid	NA	
Matrix	SEDIMENT	
Sample Size	11.03	
Size Unit-Basis	G DRY	
Units	NG/G_DRY	
C1-Chrysenes	U	
C2-Chrysenes	Ŭ	
C3-Chrysenes	Ŭ	
C4-Chrysenes	Ŭ	
Benzo(b)fluoranthene	Ũ	
Benzo(k)fluoranthene	Ŭ	
Benzo(e)pyrene	Û	
Benzo(a)pyrene	U	
Perylene	Û	
ndeno(1,2,3-cd)pyrene	Û	
Dibenz(a,h)anthracene	U	
Benzo(g,h,i)perylene	Ū	
Total PAH	0.94 J	
Surrogate Recoveries (%)		
Naphthalene-d8	108	
Acenaphthene-d10	96	
Phenanthrene-d10	93	
Benzo(a)pyrene-d12	121 N	
bb(H)-Cholane	100	

The Business of Innovation

Client ID	060208-03: Sand White Quartz, -50+7(
Part In the						
Battelle ID	BL034LCS-F					
Sample Type	LCS					
Collection Date	10/15/07					
Extraction Date	10/15/07					
Analysis Date	10/24/07					
Analytical Instrument	MS					
% Moisture	NA					
% Lipid	NA					
Matrix	SEDIMENT					
Sample Size	NA					
Size Unit-Basis	NA					
Units	NG	1	Target	% Recovery	Qualifier	
C3-Alkylbenzenes		U				
C4-Alkylbenzenes		Ŭ				
C5-Alkylbenzenes		ŭ				
C6-Alkylbenzenes		ŭ				
Benzo(b)thiophene	2229.17		2508.50	89		
C1-benzo(b)thiophenes		U	2000.00	00		
C2-benzo(b)thiophenes		ŭ				
C3-benzo(b)thiophenes		Ŭ				
C4-benzo(b)thiophenes		Ũ				
Naphthalene	2498.65		2500.50	100		
C1-Naphthalenes	100001	U		100		
C2-Naphthalenes		ū				
C3-Naphthalenes		Ū				
C4-Naphthalenes		U				
C1-Biphenyls + Dibenzofuran		Ú				
Biphenyl	2082.17		2504.25	83		
C2-Biphenyls + C1-Dibenzofurans		U	2007.00			
Acenaphthylene	2301.46		2502.25	92		
Acenaphthene	2451.23		2501.63	98		
Dibenzofuran	2068.13		2504.75	83		
Fluorene	2285.22		2501.38	91		
C1-Fluorenes		U	Decelores.			
C2-Fluorenes		U				
C3-Fluorenes		U				
C4-Fluorenes		U				
Anthracene	2095.12		2500.38	84		
Phenanthrene	2091		2501.25	84		
C1-Phenanthrenes/Anthracenes		U				
C2-Phenanthrenes/Anthracenes		U				
C3-Phenanthrenes/Anthracenes		U				
C4-Phenanthrenes/Anthracenes		U				
Retene		U				
Dibenzothiophene	1838.31		2502.50	73		
C1-Dibenzothlophenes		U				
C2-Dibenzothiophenes		U				
C3-Dibenzothiophenes		U				
C4-Dibenzothiophenes		U				
Fluoranthene	2132.27		2501.25	85		
Pyrene	2165.99		2500.88	87		
C1-Fluoranthenes/Pyrenes		U				
C2-Fluoranthenes/Pyrenes		U				
C3-Fluoranthenes/Pyrenes		U				
C4-Fluoranthenes/Pyrenes		U				
C0-Benzo(b)naphthothiophenes		U				
C1-Benzo(b)naphthothiophenes		U				
C2-Benzo(b)naphthothiophenes		U				
C3-Benzo(b)naphthothiophenes C4-Benzo(b)naphthothiophenes		U				
Contraction of the second s		U	- interests	1.12		
Benzo(a)anthracene	2291.31		2500.63	92		

The Business of Innovation

Client ID	060208-03: Sand, White Quartz, -50+70					
Battelle ID	BL034LCS-P					
Sample Type	LCS					
Collection Date	10/15/07					
Extraction Date	10/15/07					
Analysis Date	10/24/07					
Analytical Instrument	MS					
% Moisture	NA					
% Lipid	NA					
Matrix	SEDIMENT					
Sample Size	NA					
Size Unit-Basis	NA					
Units	NG	-	Target %	Recovery	Qualifier	
Chrysene	2168.08		2501.00	87		
C1-Chrysenes		U	202 (202	200		
C2-Chrysenes		U				
C3-Chrysenes		U				
C4-Chrysenes		U				
Benzo(b)fluoranthene	2605.55		2502.13	104		
Benzo(k)fluoranthene	2769.68		2501.38	111		
Benzo(e)pyrene	2446.43		2503.50	98		
Benzo(a)pyrene	2764.71		2502.00	111		
Perylene	2683.02		2505.13	107		
ndeno(1,2,3-cd)pyrene	2979.41		2501.25	119		
Dibenz(a,h)anthracene	2790.14		2501.38	112		
Benzo(g,h,i)perylene	2908.59		2500.75	116		
Total PAH	50416.47					
Surrogate Recoveries (%)						
Naphthalene-d8	92					
Acenaphthene-d10	92 81					
henanthrene-d10	79					
lenzo(a)pyrene-d12	111					
b(H)-Cholane	82					

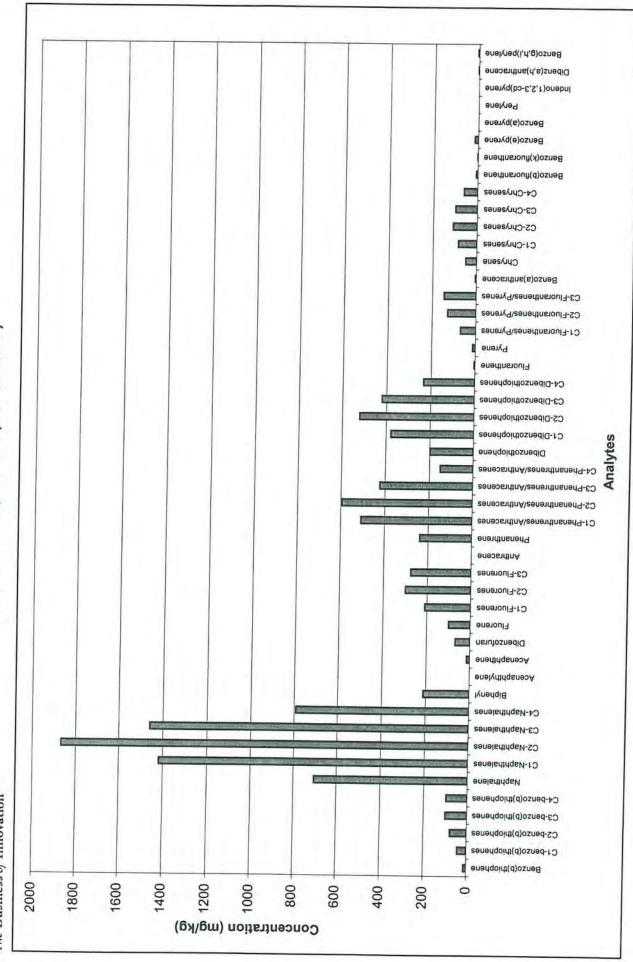
Battelle The Business of Innovation

Procedural Blank (BL033PB-P)

			C1-Chrysenes C2-Chrysenes Benzo(b)fluoranthenes
			C2-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes Benzo(a)anthracene Chrysene
			C3-Dibenzothiophenes C4-Dibenzothiophenes Pyrene C1-Fluoranthenes/Pyrenes
			C4-PhenanthrenestAnthrecenes Dibenzothiophenes C2-Dibenzothiophenes
			C3-Phenanthrenes/Anthracenes C2-Phenanthrenes/Anthracenes C2-Phenanthrenes/Anthracenes
			C3-Fluorenes C1-Fluorenes C3-Fluorenes
			Biphenyl Acensphithgene Acensphithene Dibenzotuen
			C4-Naphthalenes C3-Naphthalenes C2-Naphthalenes
			C3-Naphthalenes
	1		Naphthalene

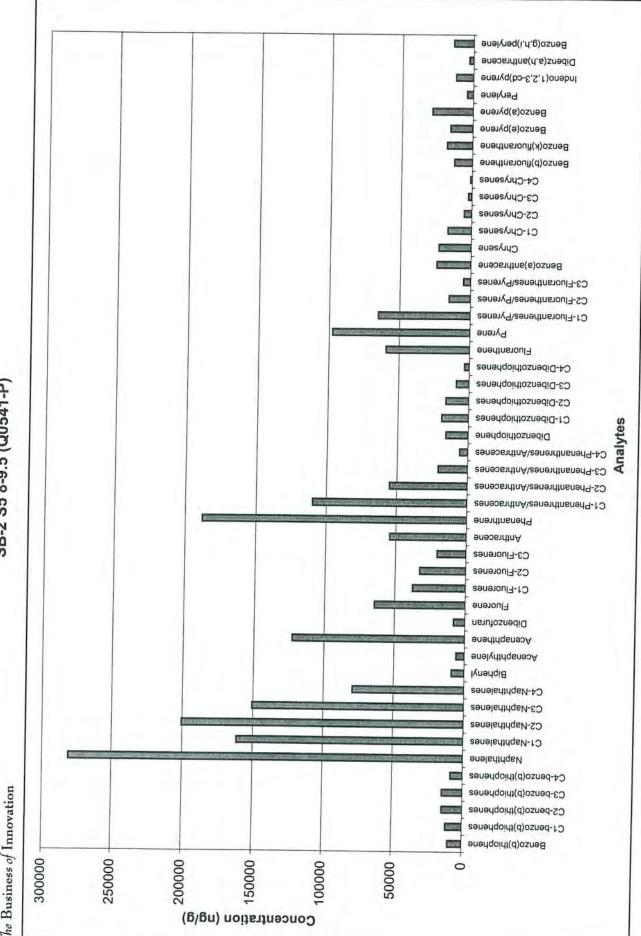


GO98: North Slope Crude (BL042NSC-P)





SB-2 S5 8-9.5 (Q0541-P)



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GP-12 S2 23-24 (Q0545-P)

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Client ID	SB-2 S5 8-9.5	GP-12 S2 23-24
Battelle ID	Q0541-P	00545.0
Sample Type		Q0545-P
Collection Date	SA	SA
Extraction Date	09/17/07	09/20/07
	10/15/07	10/15/07
Analysis Date	10/24/07	10/24/07
Analytical Instrument	MS	MS
% Moisture	18.05	26.35
% Lipid	NA	NA
Matrix	SOIL	SOIL
Sample Size	4.59	3.93
Size Unit-Basis	G DRY	G DRY
Units	NG/G_DRY	NG/G_DRY
C23 Tricyclic Terpane	808.38	U
C29 Tricyclic Terpane -22S	186.44	U
C29 Tricyclic Terpane -22R	161.44	
18a(H)-22,29,30-Trisnomeohopane -TS	106.17	U
17a(H)-22,29,30-Trisnorhopane -TM		U
30-Norhopane	206.65	9.32 J
18a(H) & 18b(H)-Oleananes	585.14	17.21 J
	249.57	Ŭ
Hopane	1145.01	25.49
30-Homohopane -22S	348.58	7.89 J
30-Homohopane -22R	256.76	9.62 J
13b(H),17a(H)-20S-Diacholestane	933.64	U
13b(H),17a(H)-20R-Diacholestane	506,1	U
14a(H),17a(H)-20R-methylcholestane	2052.28	1.56 J
14a(H),17a(H)-20S-Ethylcholestane	792.41	1.33 J
14a(H),17a(H)-20R-Ethylcholestane	1253.6	U
C21-TAS	233.87	96.2
C26-TAS(20S)	316.84	U
C26,C27-TAS	1112.01	3.93 J
C27-TAS(20R)	582.47	1.57 J
C28-TAS(20S)	356.92	1.69 J
C28-TAS(20R)	285.06	
C21-MAS	181.17	1.48 J
C22-MAS	79.43	2.44 J
C27-MAS		U
C27-20R-MAS	77.57	5.44 J
C27-20S-MAS	450.42	67.48
C28-20S-MAS	331.86	18.33
C27-C2920S/R-MAS	473.27	5.27 J
	595.97	237.34
C29-20S-MAS	1813.24	U
C29-20R-MAS	389.19	626.34
TAS_245	U	U
MAS_239	U	U
Surrogate Recoveries (%)		
Naphthalene-d8	123 N	103
Acenaphthene-d10	189 N	110
Phenanthrene-d10	94	93
Benzo(a)pyrene-d12	107 D	93 72 D
5b(H)-Cholane	209 N	
104 J. 40 11 11 11 1	203 14	1182 N

The Business of Innovation

Client ID	060208-03: Sand, White Quartz, -50+70		
Battelle ID	BL034LCS-P		
Sample Type			
Collection Date	LCS		
Extraction Date	10/15/07		
	10/15/07		
Analysis Date	10/24/07		
Analytical Instrument	MS		
% Moisture	NA		
% Lipid	NA		
Matrix	SEDIMENT		
Sample Size	NA		
Size Unit-Basis	NA		
Units	NG	Target % Recovery	Qualifier
CO2 Televalla Tarran	10		
C23 Tricyclic Terpane	U		
C29 Tricyclic Terpane -22S	U		
C29 Tricyclic Terpane -22R	U		
18a(H)-22,29,30-Trisnomeohopane -TS	U		
17a(H)-22,29,30-Trisnorhopane -TM	U		
30-Norhopane	Ú		
18a(H) & 18b(H)-Oleananes	Ũ		
Hopane	U		
30-Homohopane -22S	Ū		
30-Homohopane -22R	Ŭ		
13b(H),17a(H)-20S-Diacholestane	Ŭ		
13b(H),17a(H)-20R-Diacholestane	Ŭ		
14a(H),17a(H)-20R-methylcholestane	Ŭ		
14a(H),17a(H)-20S-Ethylcholestane	Ŭ		
14a(H),17a(H)-20R-Ethylcholestane	U		
C21-TAS	Ŭ		
C26-TAS(20S)			
C26,C27-TAS	U		
C27-TAS(20R)	U		
C28-TAS(208)	U		
C28-TAS(20S)	U		
	U		
C21-MAS	U		
C22-MAS	U		
C27-MAS	U		
C27-20R-MAS	U		
C27-20S-MAS	U		
C28-20S-MAS	U		
C27-C2920S/R-MAS	U		
C29-20S-MAS	U		
C29-20R-MAS	Ŭ		
TAS_245	U		
MAS_239	U		
Surrogate Recoveries (%)			
Naphthalene-d8	92		
cenaphthene-d10	81		
Phenanthrene-d10			
Benzo(a)pyrene-d12	79		
b(H)-Cholane	111		
and a shorane	82		

The Business of Innovation

110/0011001.1001001-0001		
Client ID	Procedural Blank	
	and a second second	
Battelle ID	BL033PB-P	
Sample Type	PB	
Collection Date	10/15/07	
Extraction Date	10/15/07	
Analysis Date	10/24/07	
Analytical Instrument	MS	
% Moisture	22.07	
% Lipid	NA	
Matrix	SEDIMENT	
Sample Size		
	11.03	
Size Unit-Basis	G_DRY	
Units	NG/G_DRY	
C23 Tricyclic Terpane	11	
C29 Tricyclic Terpane -22S	U	
	U	
C29 Tricyclic Terpane -22R	U	
18a(H)-22,29,30-Trisnomeohopane -TS	U	
17a(H)-22,29,30-Trisnorhopane -TM	U	
30-Norhopane	U	
18a(H) & 18b(H)-Oleananes	U	
Hopane	U	
30-Homohopane -22S	Ŭ	
30-Homohopane -22R	Ŭ	
13b(H),17a(H)-20S-Diacholestane	Ŭ	
13b(H),17a(H)-20R-Diacholestane	Ŭ	
14a(H),17a(H)-20R-methylcholestane		
14a(H) 17a(H) 200 Ethylohalastane	U	
14a(H),17a(H)-20S-Ethylcholestane	U.	
14a(H),17a(H)-20R-Ethylcholestane	U	
C21-TAS	U	
C26-TAS(20S)	U	
C26,C27-TAS	U	
C27-TAS(20R)	0.23 J	
C28-TAS(20S)	0.16 J	
C28-TAS(20R)	U	
C21-MAS	Ŭ	
C22-MAS	Ŭ	
C27-MAS	Ŭ	
C27-20R-MAS	Ŭ	
C27-20S-MAS	Ŭ	
C28-20S-MAS		
C27-C2920S/R-MAS	0	
C29-20S-MAS	U	
	U	
C29-20R-MAS	U U	
TAS_245	U	
MAS_239	U	
Surrogate Recoveries (%)		
Naphthalene-d8	105	
	108	
Acenaphthene-d10	96	
Phenanthrene-d10	93	
Benzo(a)pyrene-d12	121 N	
5b(H)-Cholane	100	

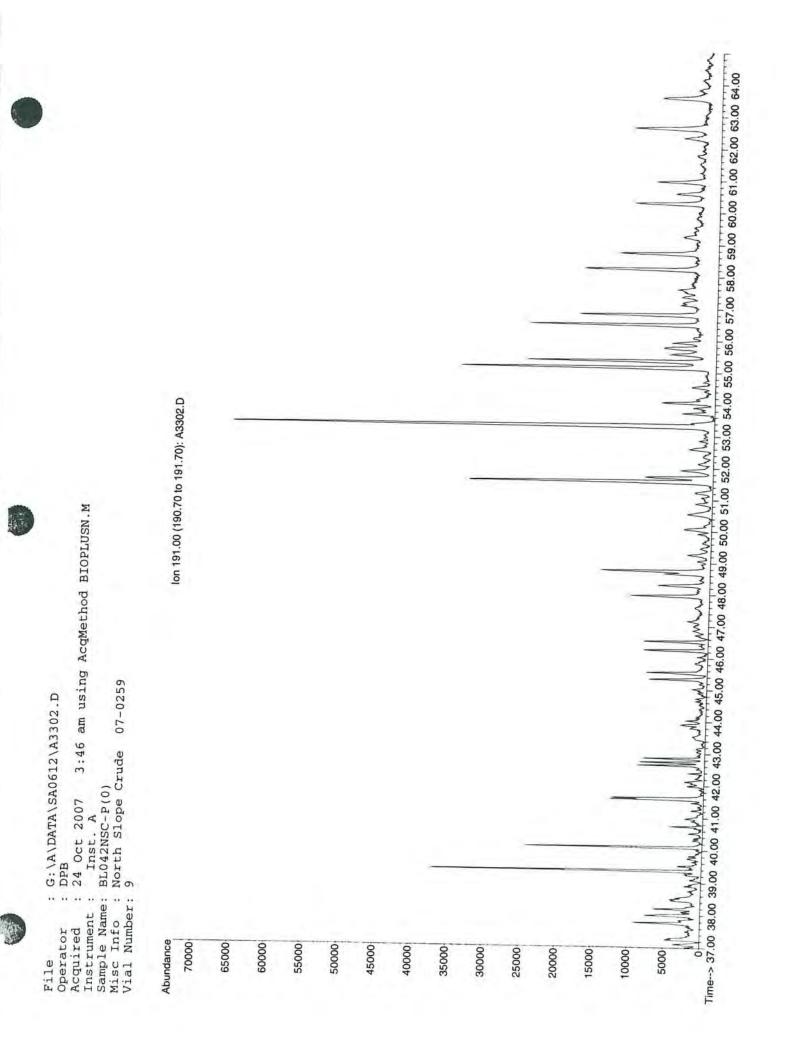
The Business of Innovation

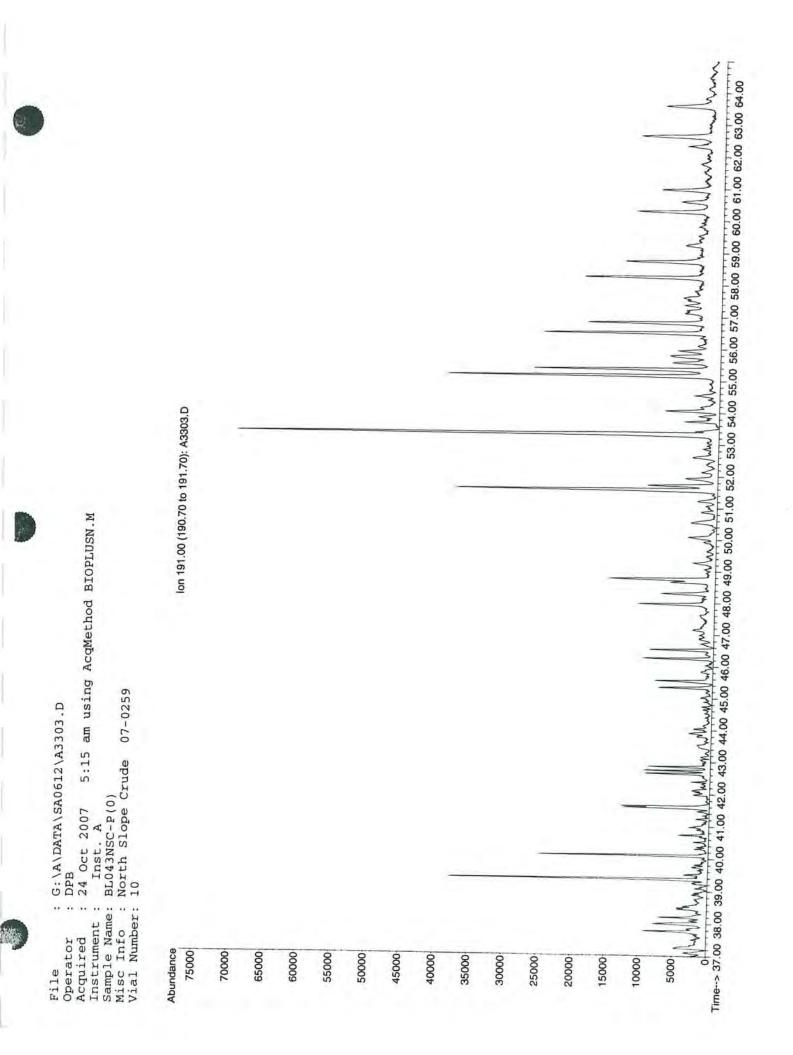
Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

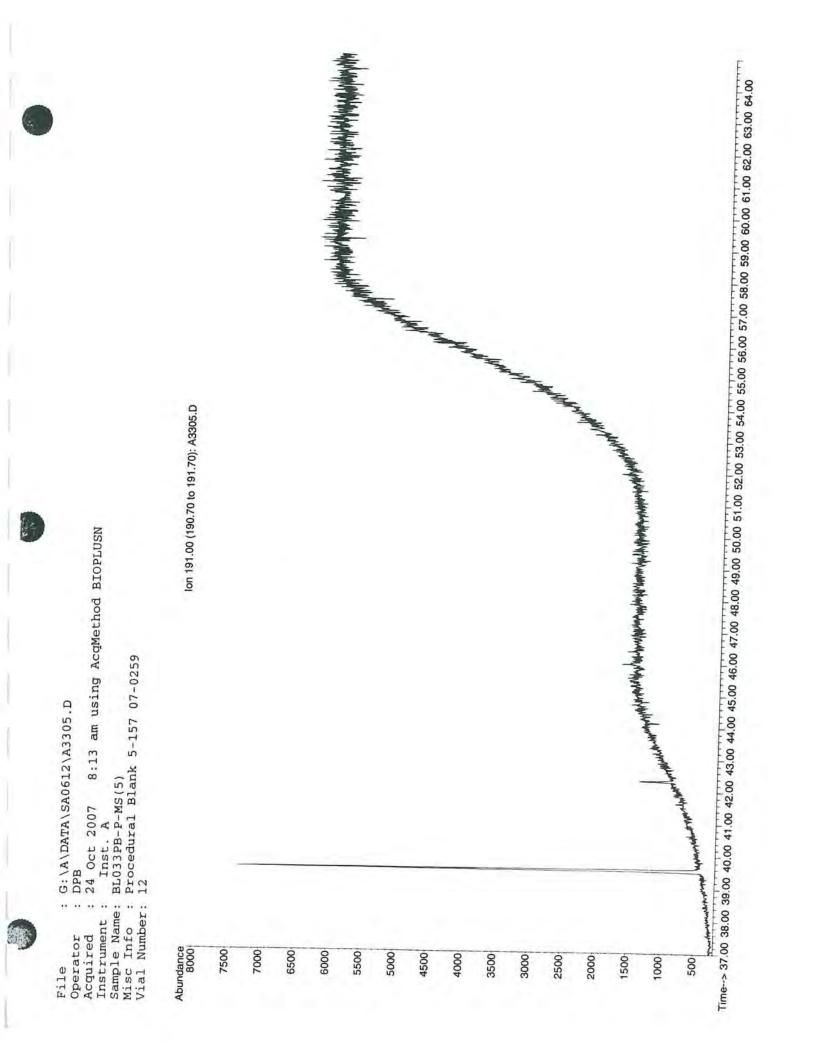
Client ID	GO98: North Slope Crude			
Detter ID	Contractor In			
Battelle ID	BL043NSC-P			
Sample Type	NSC			
Collection Date	10/17/2007			
Extraction Date	10/17/2007			
Analysis Date	10/24/2007			
Analytical Instrument	MS			
% Moisture	NA			
% Lipid	NA			
Matrix	OIL			
Sample Size	5.04			
Size Unit-Basis	MG OIL			
Units	MG/KG_OIL	Target % I	Difference	Qualifie
C22 Triquelle Terrere				
C23 Tricyclic Terpane	38.41	47.76	19.6	
C29 Tricyclic Terpane -22S	12.22	14.70	16.9	
C29 Tricyclic Terpane -22R	11.89	14.64	18.8	
18a(H)-22,29,30-Trisnomeohopane -TS	14.61	15.96	8.5	
17a(H)-22,29,30-Trisnorhopane -TM	23.47	24.82	5.4	
30-Norhopane	63.43	69.58	8.8	
18a(H) & 18b(H)-Oleananes	U			
Hopane	111.6	120.14	7.1	
30-Homohopane -22S	60.02	59.93	0.2	
30-Homohopane -22R	42.46	39.69	7.0	
13b(H), 17a(H)-20S-Diacholestane	38.87	44.18	12.0	
13b(H),17a(H)-20R-Diacholestane	23.96	25.52	6.1	
14a(H),17a(H)-20R-methylcholestane	31.74	33.94	6.5	
14a(H),17a(H)-20S-Ethylcholestane	37.87	35.93	5.4	
14a(H),17a(H)-20R-Ethylcholestane	38	39.17	3.0	
C21-TAS	U			
C26-TAS(20S)	U			
26,C27-TAS	U			
227-TAS(20R)	U			
C28-TAS(20S)	U			
228-TAS(20R)	U			
C21-MAS	U			
22-MAS	U			
27-MAS	U			
27-20R-MAS	U			
27-20S-MAS	U			
28-20S-MAS	U			
27-C2920S/R-MAS	u			
29-20S-MAS	U			
29-20R-MAS	U			
AS_245	U			
IAS 239	U			

Surrogate Recoveries (%)

Naphthalene-d8 Acenaphthene-d10 Phenanthrene-d10 Benzo(a)pyrene-d12 5b(H)-Cholane

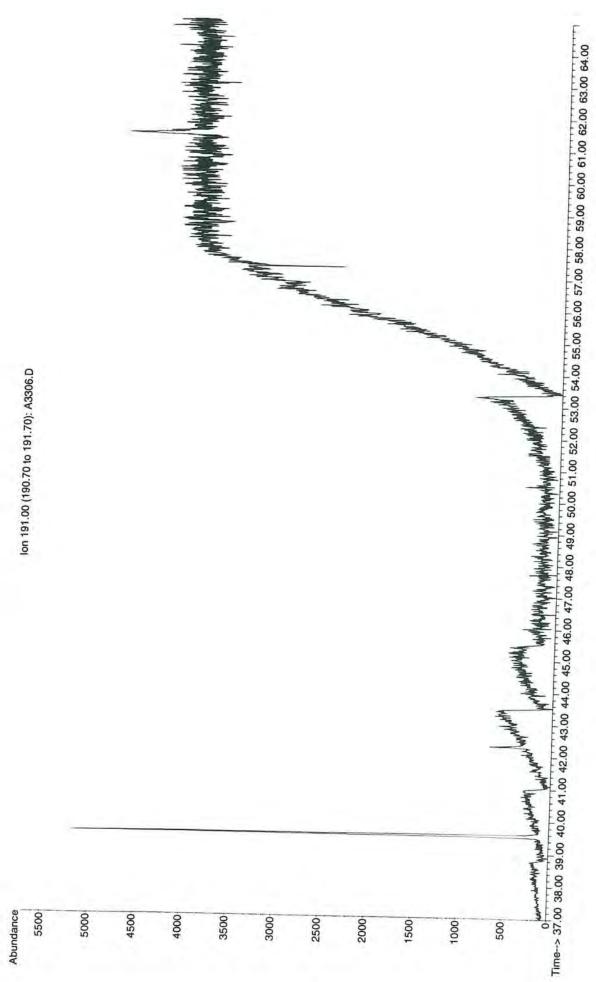


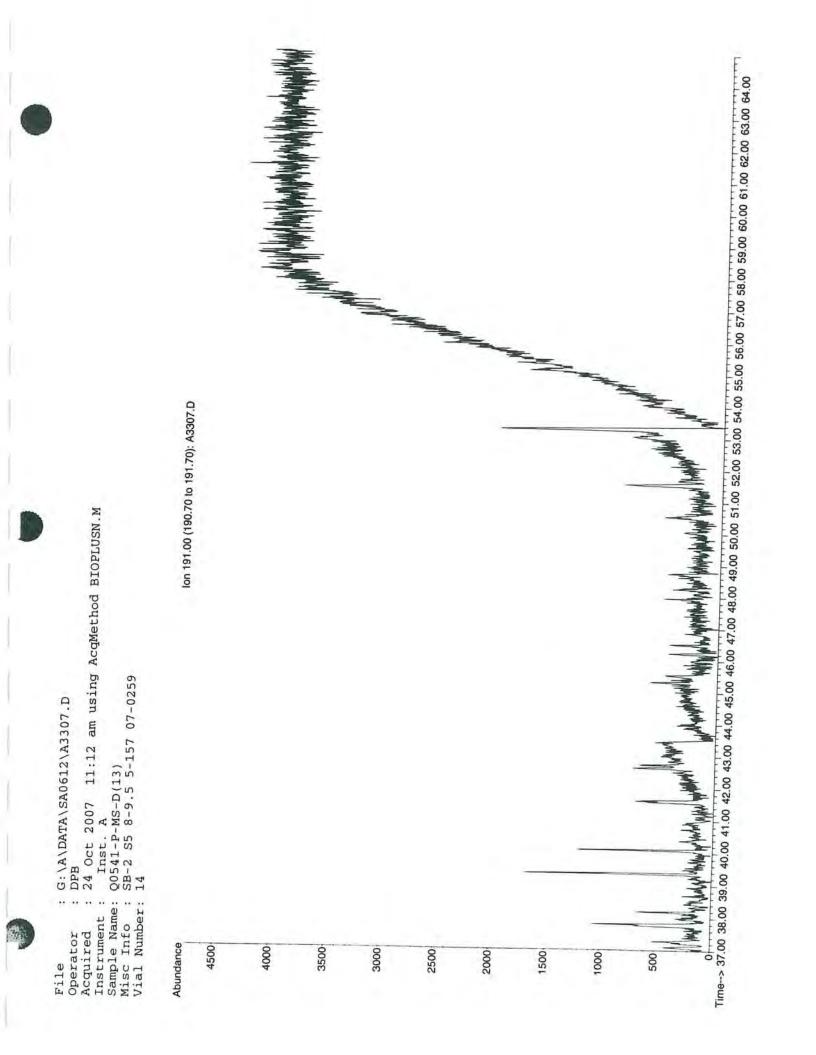


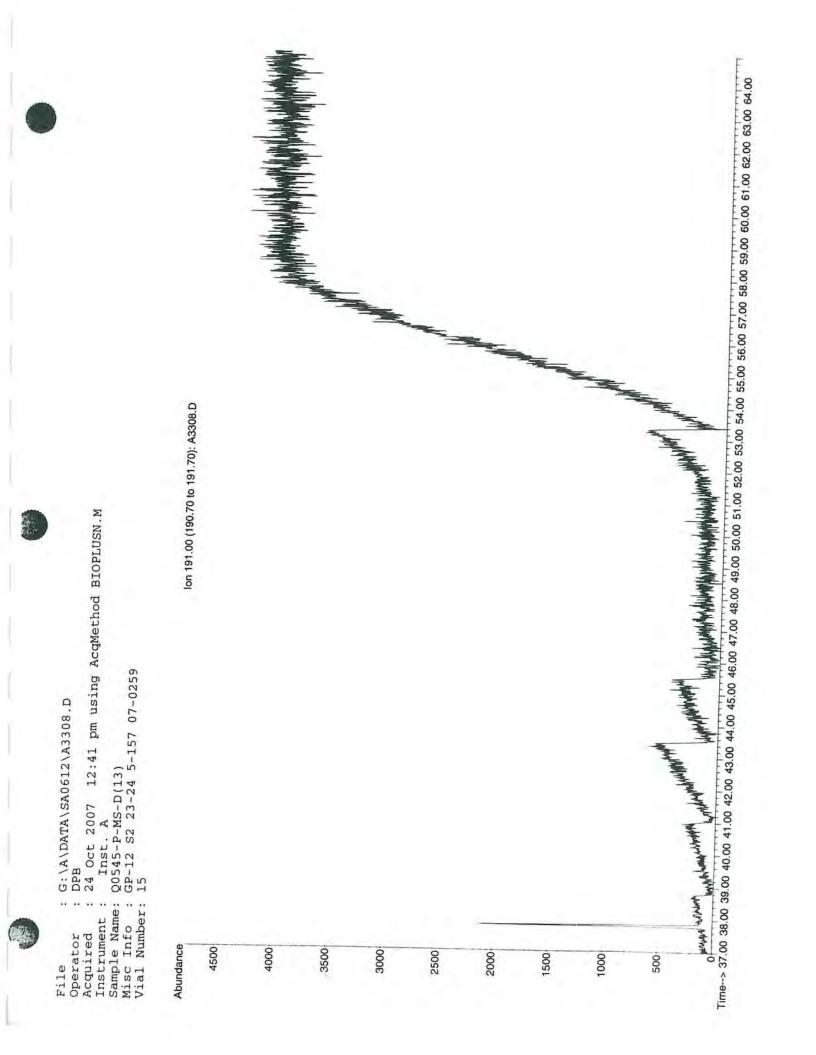


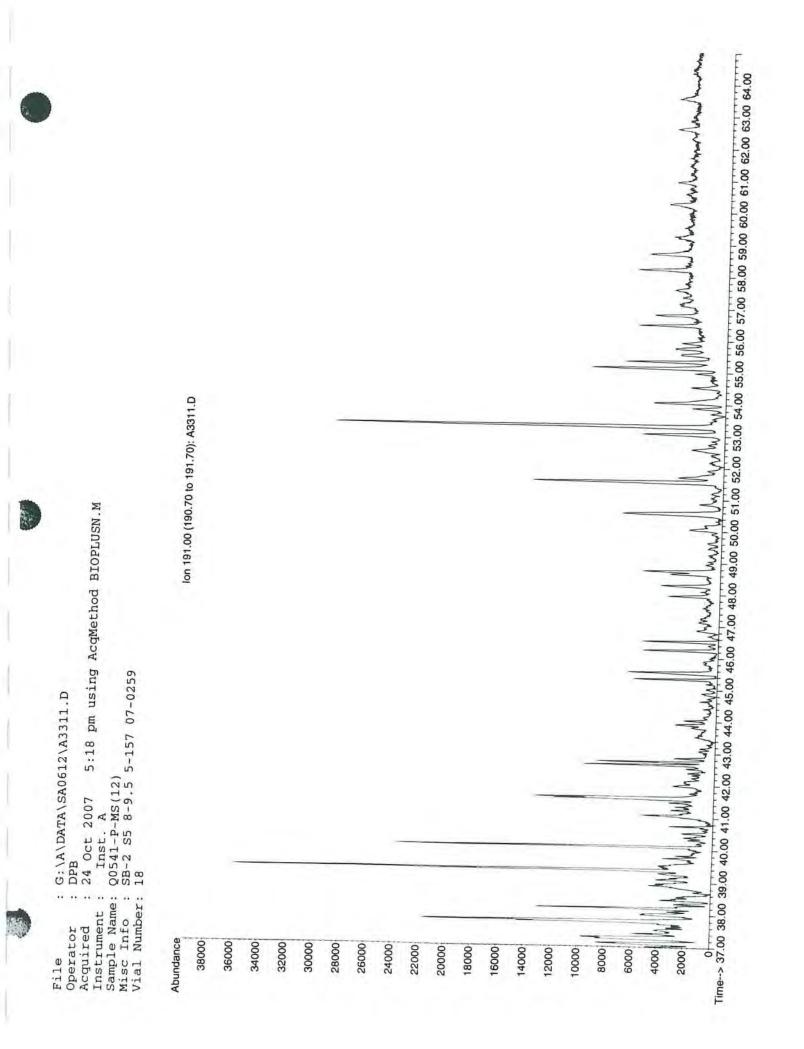


1306.D	24 Oct 2007 9:42 am using AcqMethod BIOPLUSN.M	buo34LCS-P-MS(5) Laboratory Control Sample 5-157 07-0259 13	
: G:\A\DATA\SA0612\A3306.D : DPB	24 Oct 2007 9:42 Inst. A	Multiple Number: BLU34LCS-P-MS(5) Misc Info : Laboratory Control Mal Number: 13	
File : Operator :	scquired : Instrument :	Wisc Info : Vial Number:	









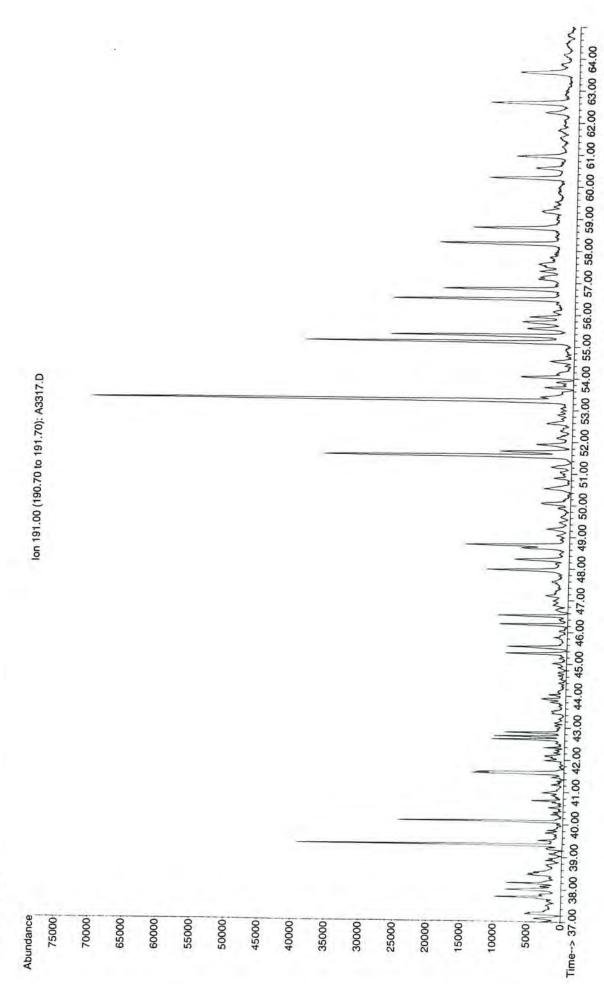


AcqMethod BIOPLUSN.M	
G:\A\DATA\SA0612\A3312.D DPB 24 Oct 2007 6:47 pm using AcqMethod BIOPLUSN.M Inst. A Q0545-P-MS(12) GP-12 S2 23-24 5-157 07-0259 19	
File :	

Abundance	100000	00006	80000	70000	60000	50000	40000	30000	20000	10000	
lon 191.00 (190.70 to 191.70): A3312.D											
Ο											

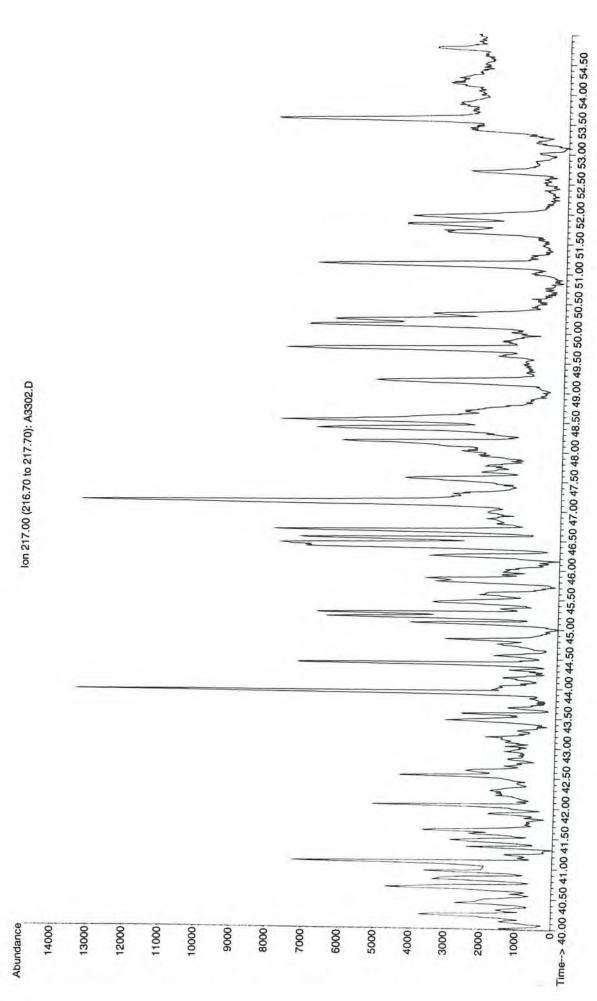


2:12 am using AcqMethod BIOPLUSN.M 07-0259 G: \A\DATA\SA0612\A3317.D Inst. A BL043NSC-P(0) North Slope Crude 25 Oct 2007 DPB Misc Info : Nor Vial Number: 24 .. Sample Name: Instrument Operator Acquired File



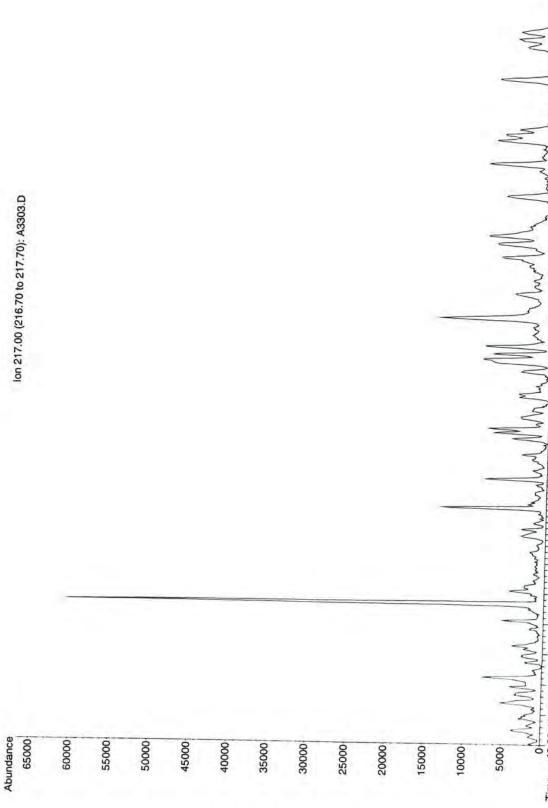


3:46 am using AcqMethod BIOPLUSN.M 07-0259 : G:\A\DATA\SA0612\A3302.D : DPB wistrument : Inst. A Sample Name: BL042NSC-P(0) Misc Info : North Slope Crude Vial Number: 9 24 Oct 2007 **Operator** Acquired File





Acquired : 24 Oct 2007 5:15 am using AcqMethod BIOPLUSN.M Instrument : Inst. A Sample Name: BL043NSC-P(0) Misc Info : North Slope Crude 07-0259 Vial Number: 10	ii.	D. 2022A	
Misc Info : North Slope Crude 07-0259 Vial Number: 10	ed : 24 Oct 2007 5; nent : Inst. A Name: BL043NSC-P(0)	.5 am using	AcqMethod BIOPLUSN.M
	ıfo : North Slope Crud ımber: 10	07-0259	
Abundance			

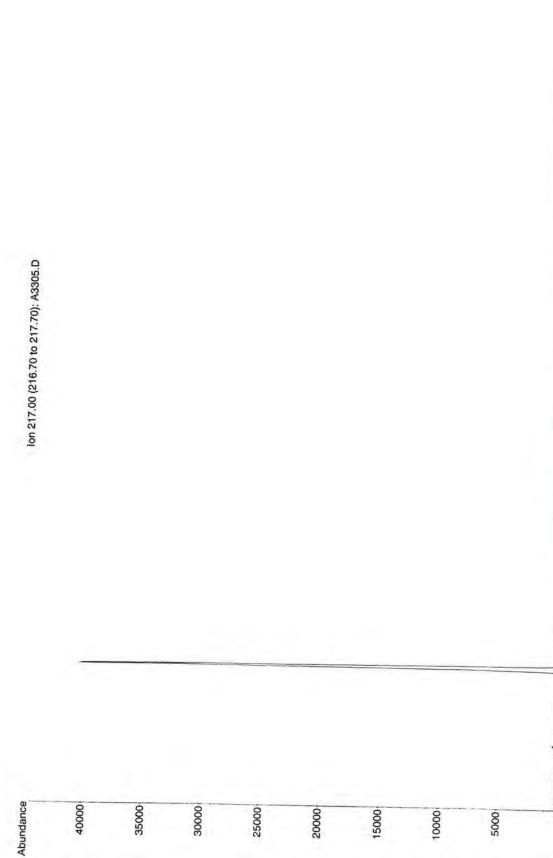


Time-> 40.00 40.50 41.00 41.50 42.50 43.50 43.50 44.50 45.50 45.50 45.50 45.50 45.50 48.50 47.50 48.50 48.50 48.50 48.50 48.50 50.50 50.50 50.50 51.50 52.00 52.50 53.50 53.50 54.50 54.50



	BIOPLI
	AcqMethod
D.D	using
305	am
612\A3	8:13
TA\SA061	2007
A\DA	Oct

NSD Instrument : Inst. A Sample Name: BL033PB-P-MS(5) Misc Info : Procedural Blank 5-157 07-0259 Vial Number: 12 : G:\A : DPB 24 ... File Operator Acquired Instrument



Time-> 40.00 40.50 41.00 41.50 42.00 42.50 43.00 43.50 44.00 44.50 45.00 45.50 46.00 46.50 47.00 47.50 48.00 48.50 49.00 49.50 50.00 50.50 51.00 51.50 52.00 52.50 53.00 53.50 54.00 54.50

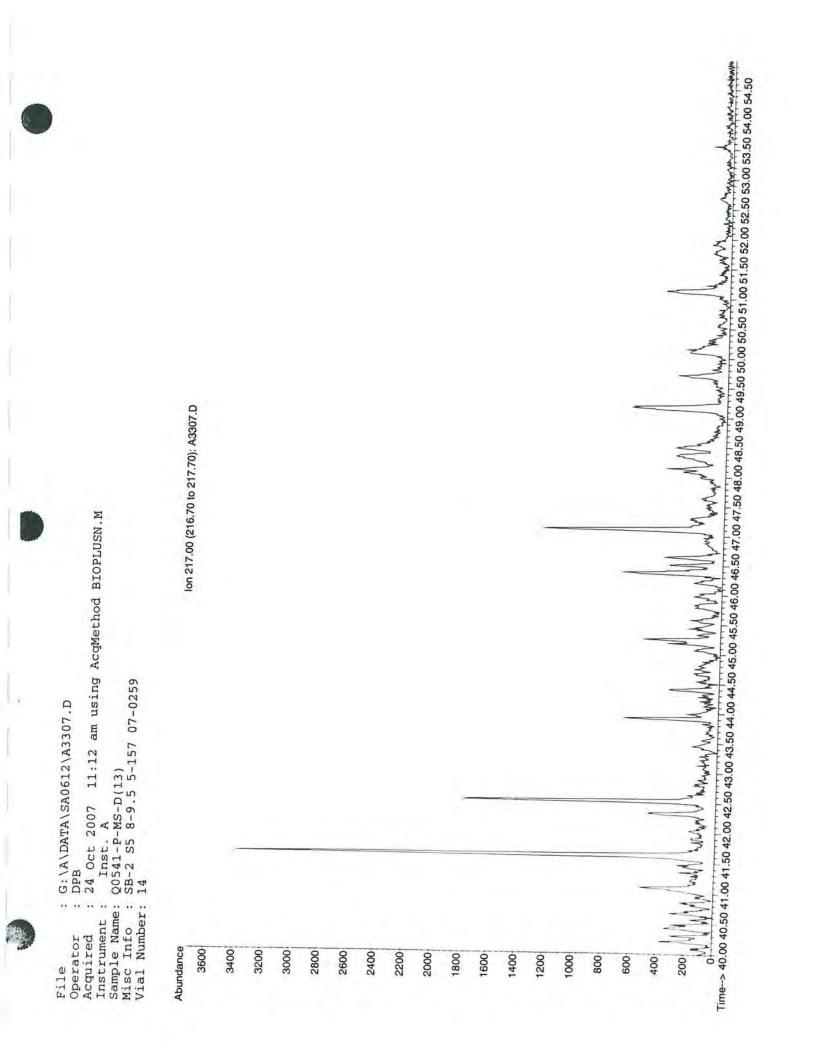


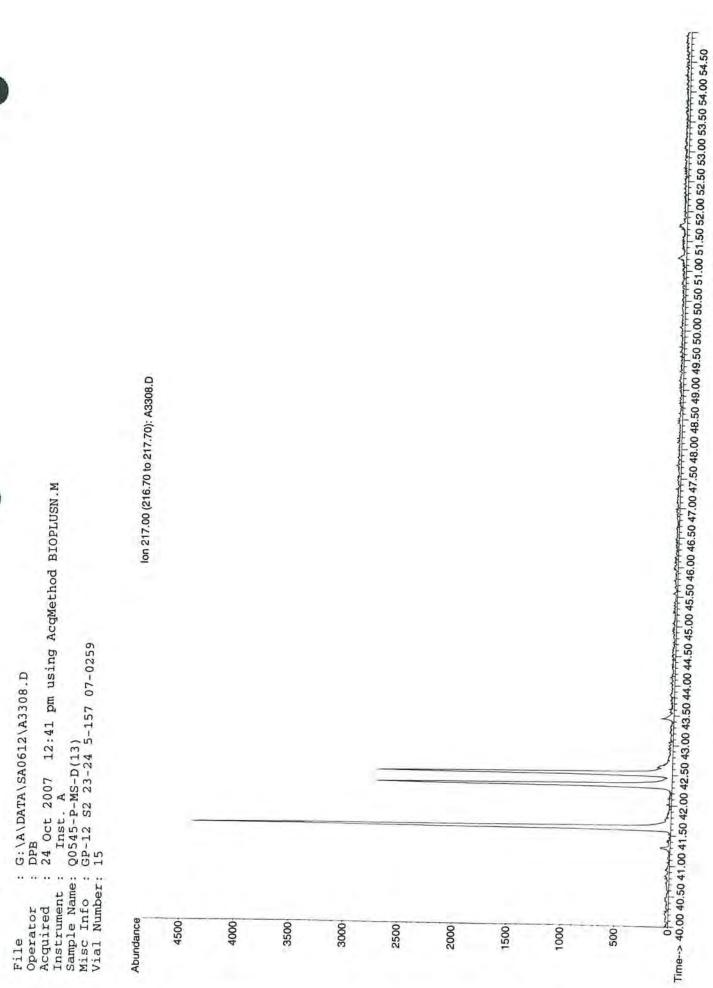
File

8	1	1	
7			

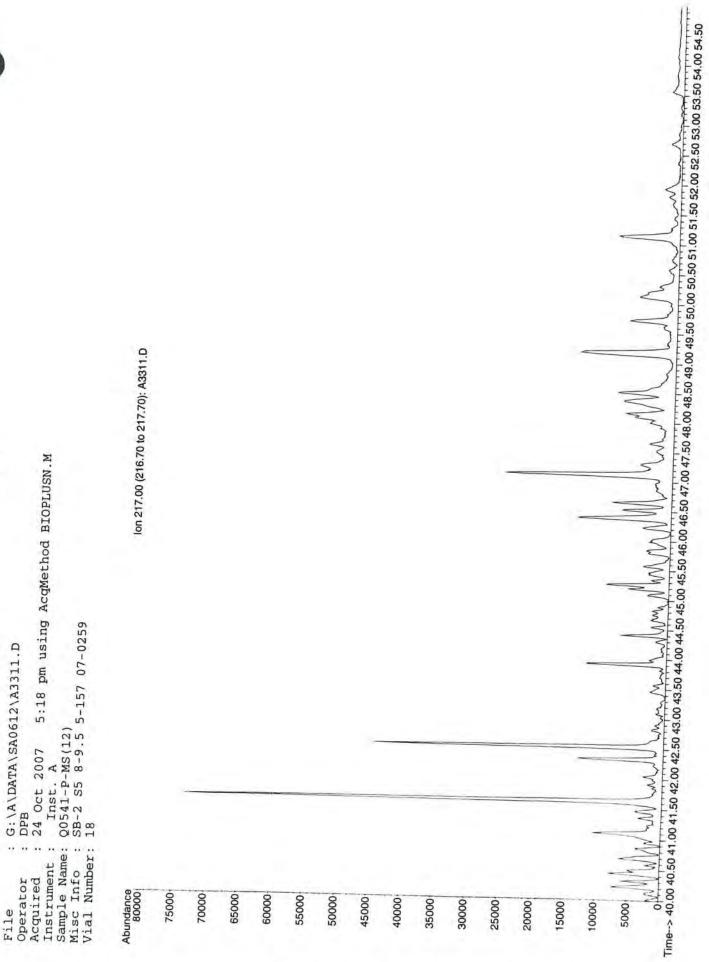
9:42 am using AcqMethod BIOPLUSN.M File : G:\A\DATA\SA0612\A3306.D Operator : DPB Acquired : 24 Oct 2007 9:42 am using AcqMethod B1 Instrument : Inst. A Sample Name: BL034LCS-P-MS(5) Misc Info : Laboratory Control Sample 5-157 07-0259 Vial Number: 13

Time--> 40.00 40.50 41.00 41.50 42.00 42.50 43.00 43.50 44.50 45.50 45.50 46.00 46.50 47.00 47.50 48.00 48.50 49.00 49.50 50.00 50.50 51.00 51.50 52.50 53.50 53.50 54.50 54.50



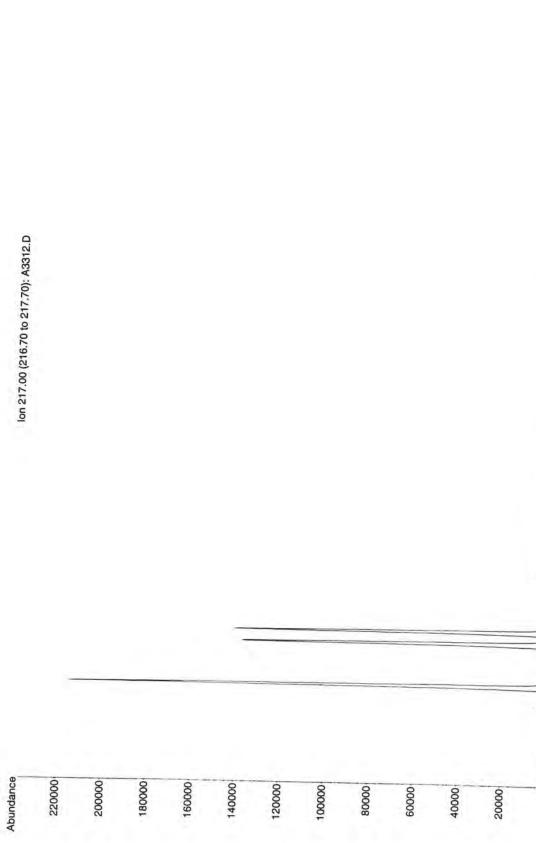




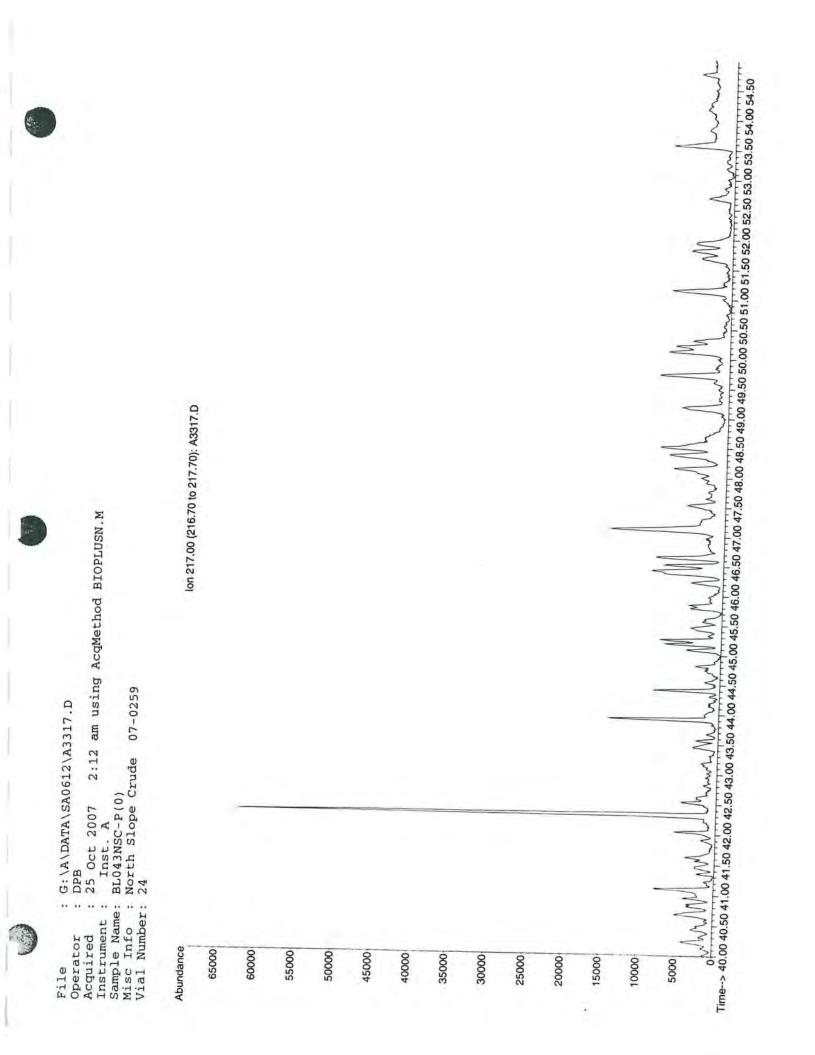




	AcqMethod BIOPLUSN.M	
: G:\A\DATA\SA0612\A3312.D : DPR	Acquired : 24 Oct 2007 6:47 pm using AcqMethod BIOPLUSN.M Instrument : Inst. A	00545-P-MS(12) GP-12 S2 23-24 5-157 07-0259 19
File : Operator :	Acquired :	sample Name: Misc Info : Vial Number:

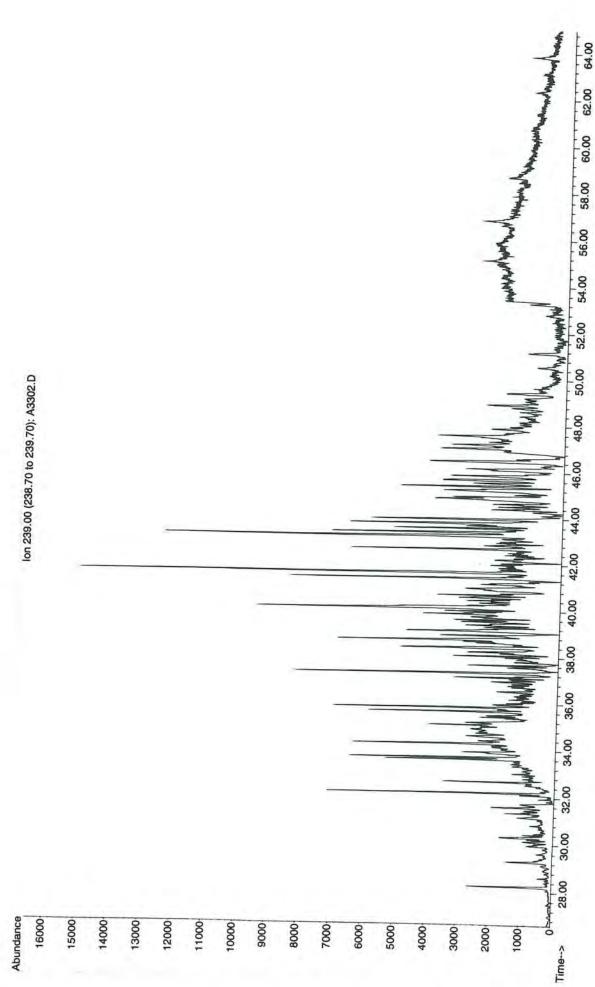


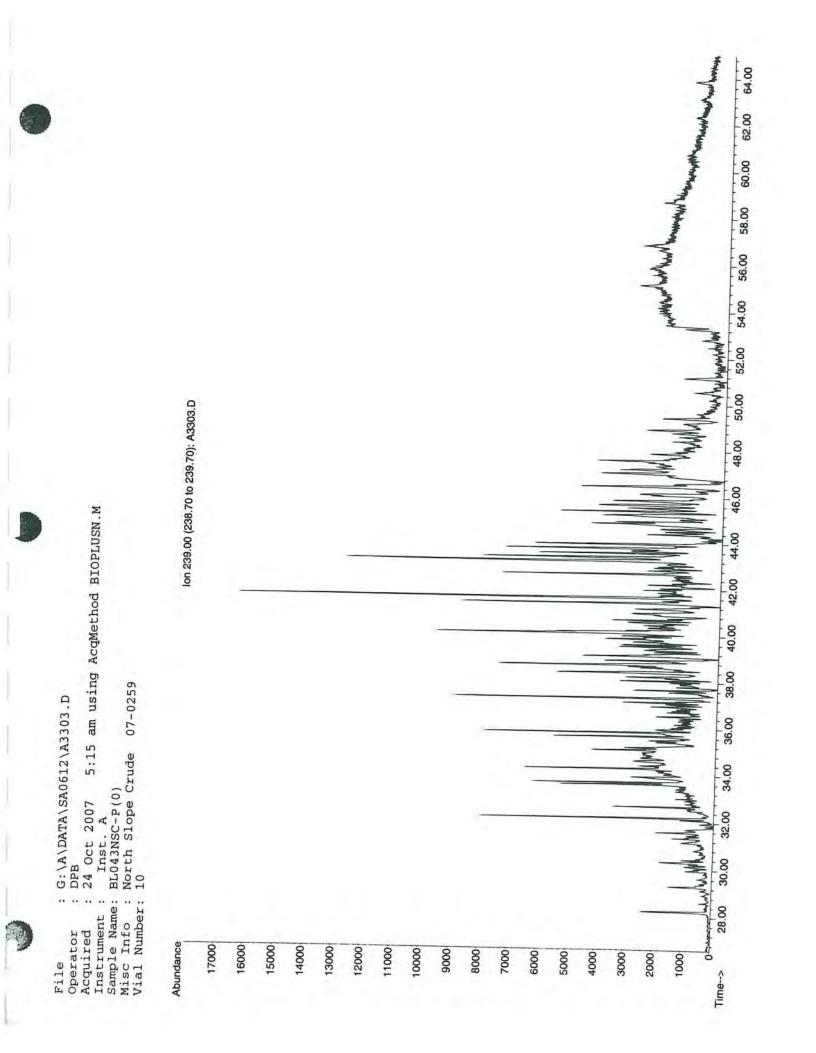
Time--> 40.00 40.50 41.00 41.50 42.00 42.50 43.00 43.50 44.00 44.50 45.00 45.50 46.00 45.50 47.00 47.50 48.00 48.50 49.00 49.50 50.00 50.50 51.00 51.50 52.00 52.50 53.00 53.50 54.00 54.50





3:46 am using AcqMethod BIOPLUSN.M 07-0259 G:\A\DATA\SA0612\A3302.D DPB : 24 Oct 2007 3:46 a : Inst. A ie: BL042NSC-P(0) : North Slope Crude 0 6 Sample Name: F Misc Info : N Vial Number: 5 Instrument File Operator Acquired

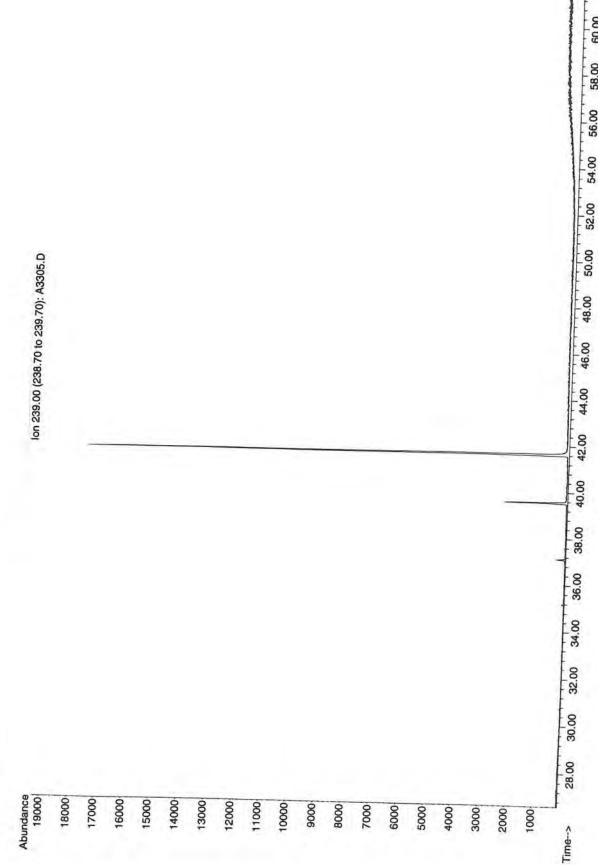








8:13 am using AcqMethod BIOPLUSN File : G:\A\DATA\SA0612\A3305.D Operator : DPB Acquired : 24 Oct 2007 8:13 am using Acq Instrument : Inst. A Sample Name: BL033PB-P-MS(5) Misc Info : Procedural Blank 5-157 07-0259 Vial Number: 12 File



64.00

62.00

60.00

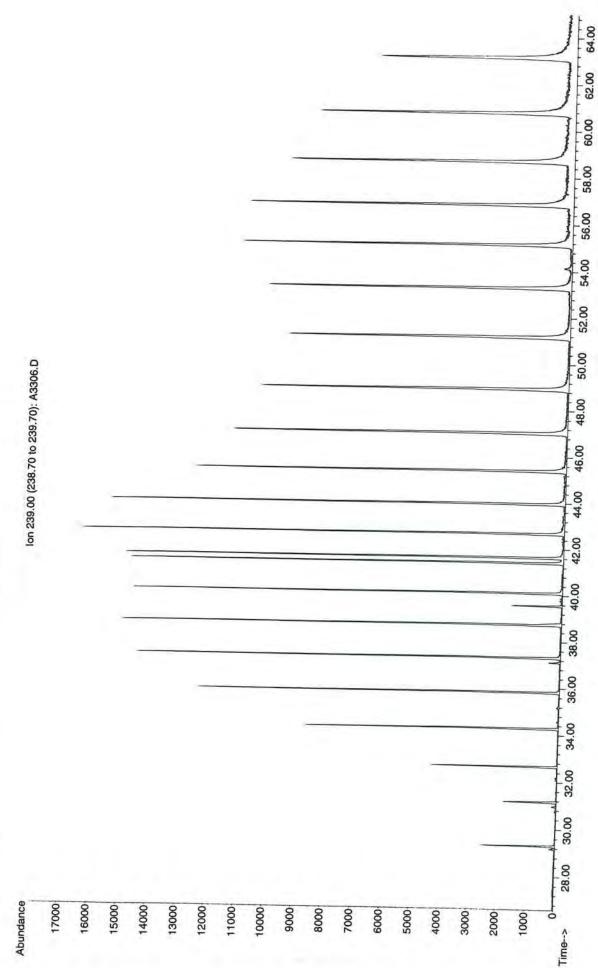
58.00

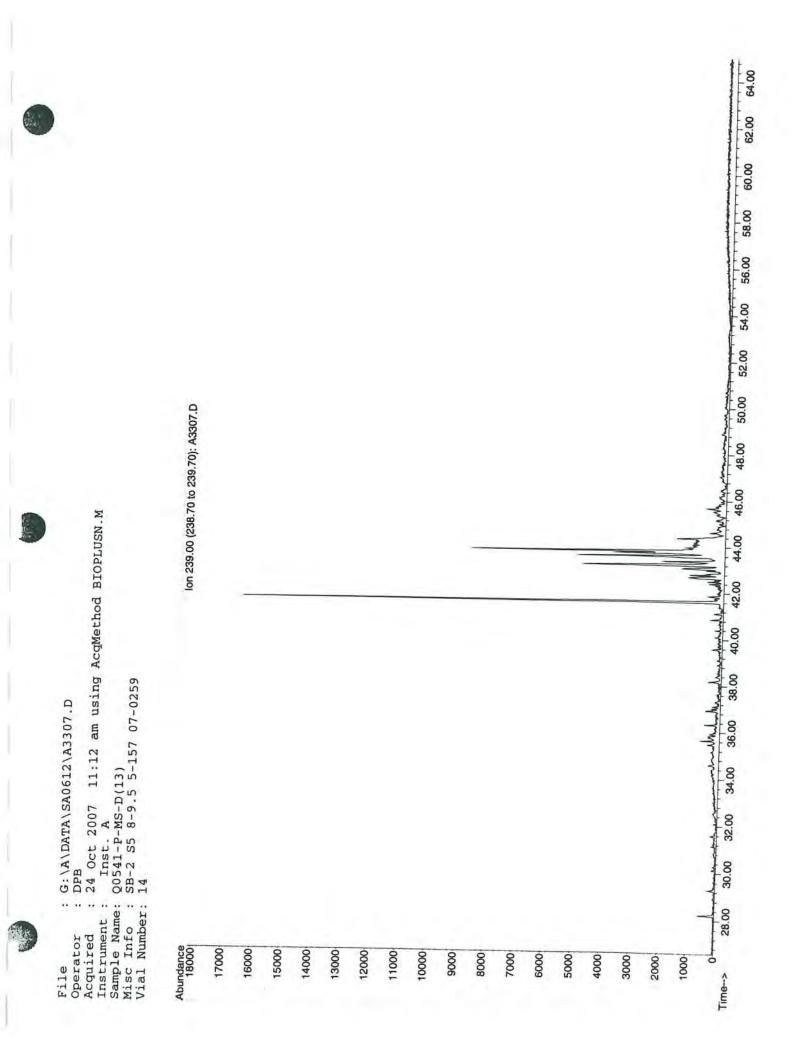
56.00

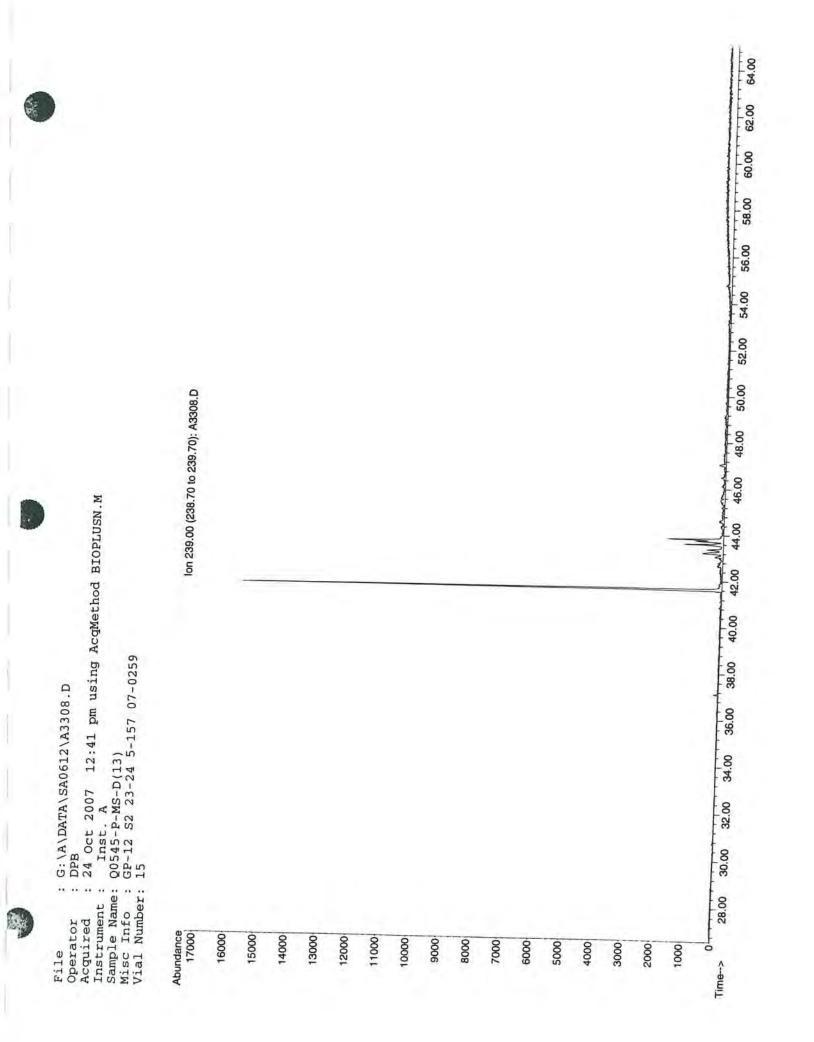


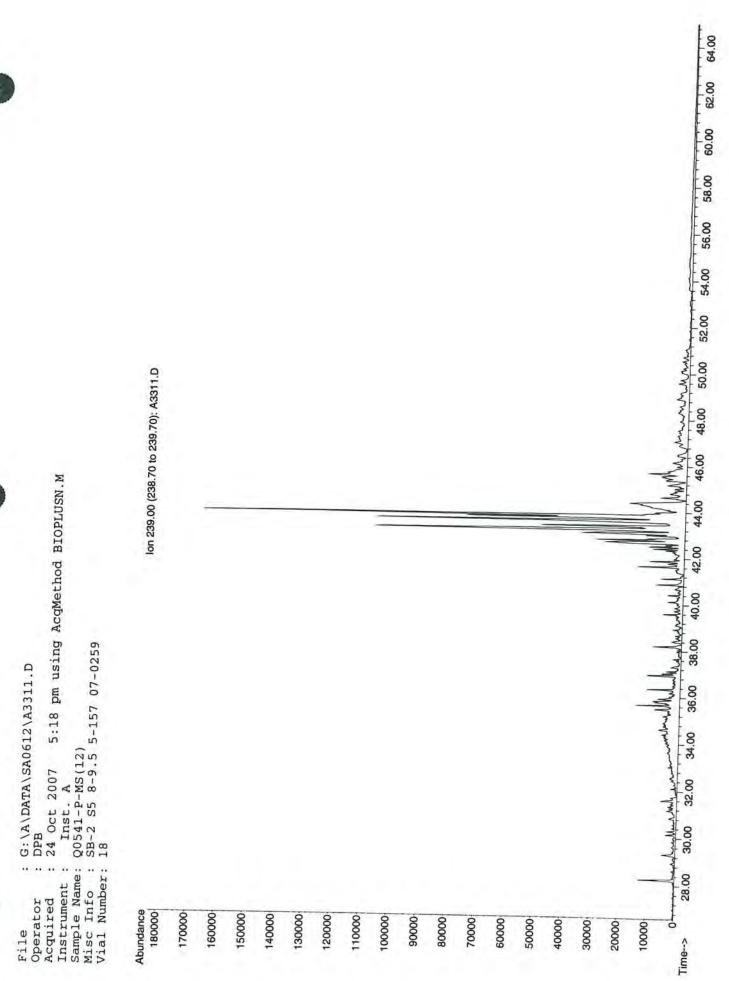


9:42 am using AcqMethod BIOPLUSN.M G:\A\DATA\SA0612\A3306.D DPB 24 Oct 2007 9:42 am using AcqMethod BI Inst. A BL034LCS-P-MS(5) Laboratory Control Sample 5-157 07-0259 13 Sample Name: F Misc Info : I Vial Number:] Instrument **Operator** Acquired File



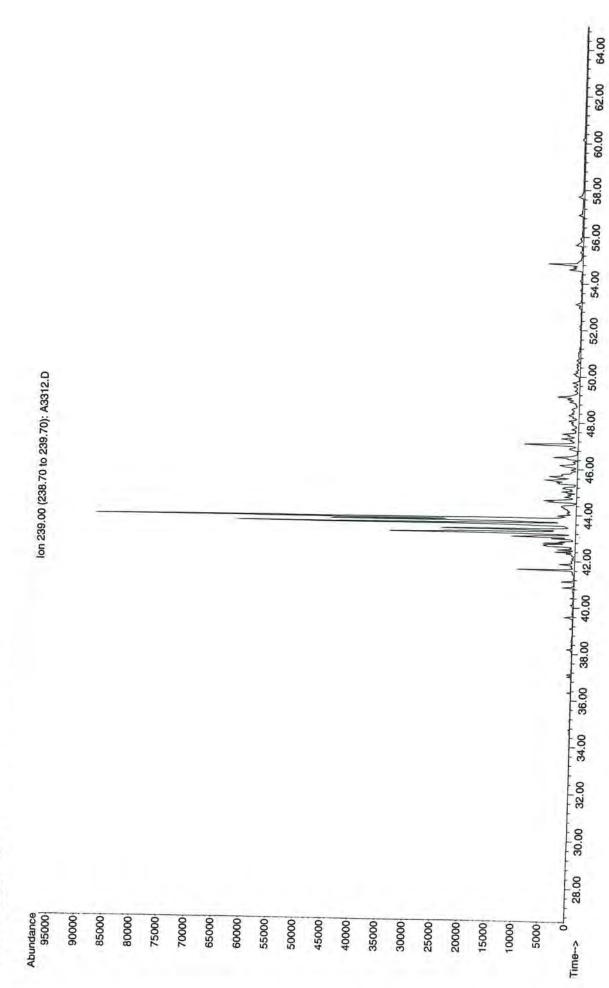








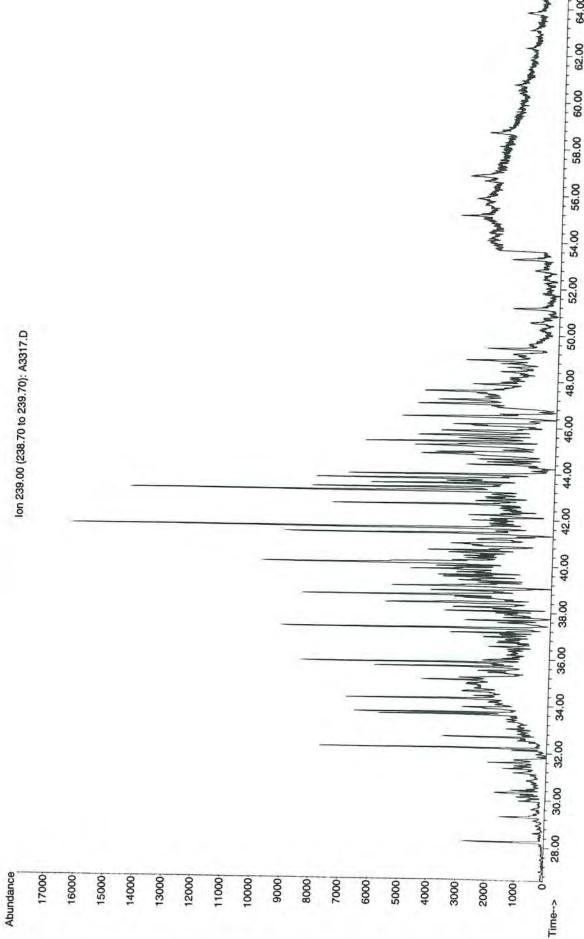
	6:47 pm using AcqMethod BIOPLUSN.M	0259
G:\A\DATA\SA0612\A3312.D	24 Oct 2007 6:47 pm us: Thet a	Q0545-P-MS(12) GP-12 S2 23-24 5-157 07-0259 19
File : Operator :	Acquired : Instrument :	Sample Name: Misc Info : Vial Number:





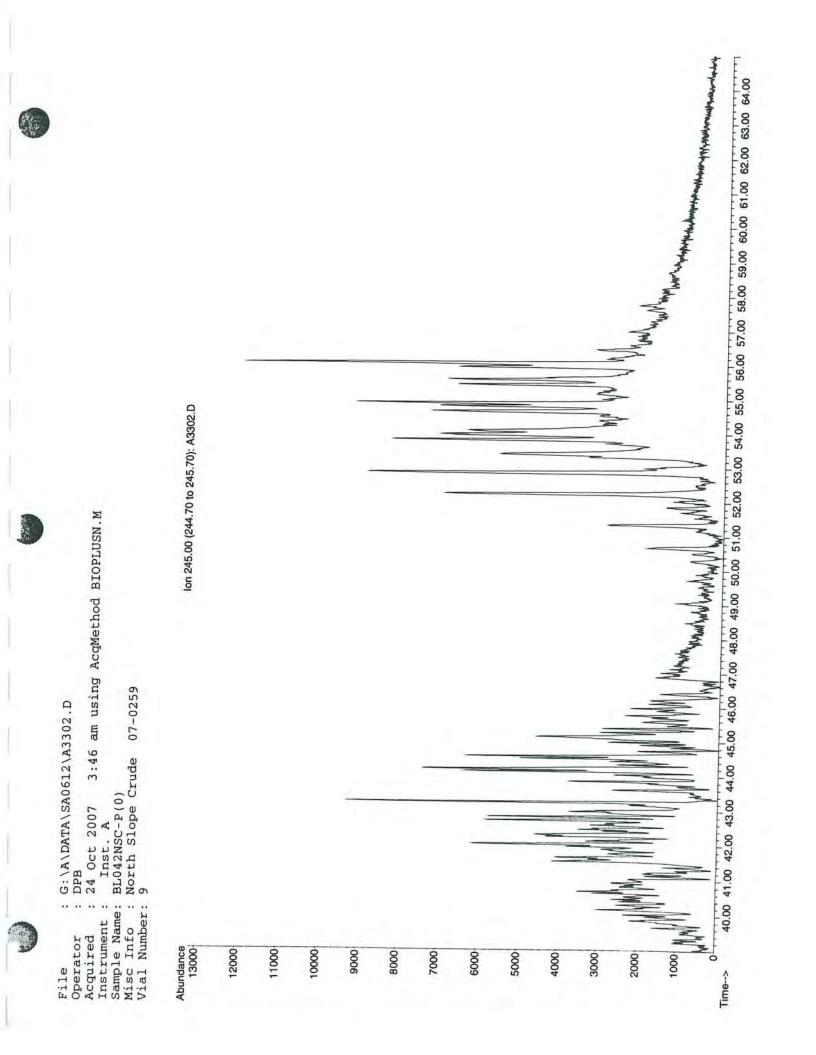
2:12 am using AcqMethod BIOPLUSN.M 07-0259 File : G:\A\DATA\SA0612\A3317.D Operator : DPB Acquired : 25 Oct 2007 2:12 am usi Instrument : Inst. A Sample Name: BL043NSC-P(0) Misc Info : North Slope Crude 07-025 Vial Number: 24





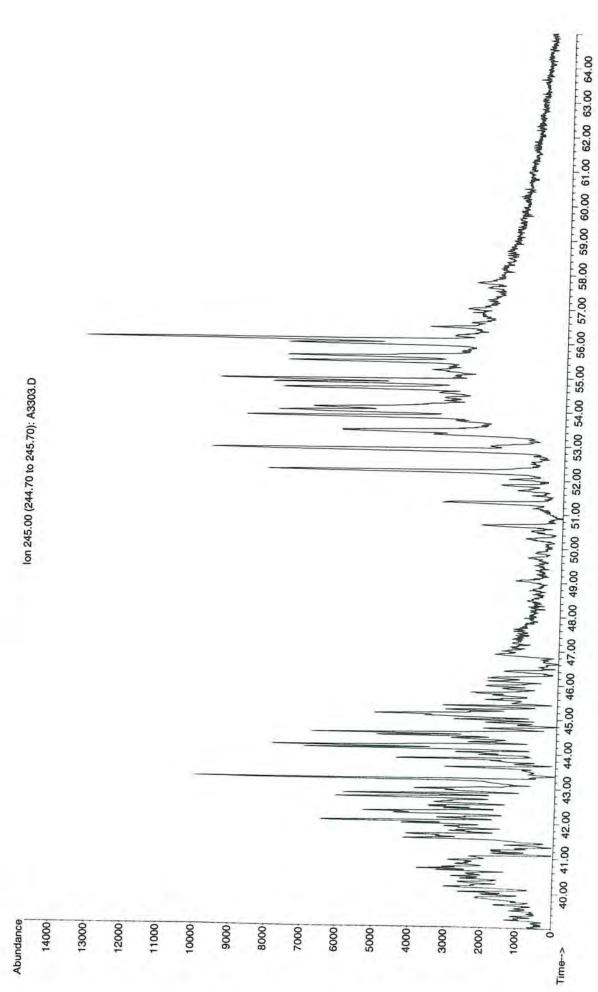
64.00

62.00



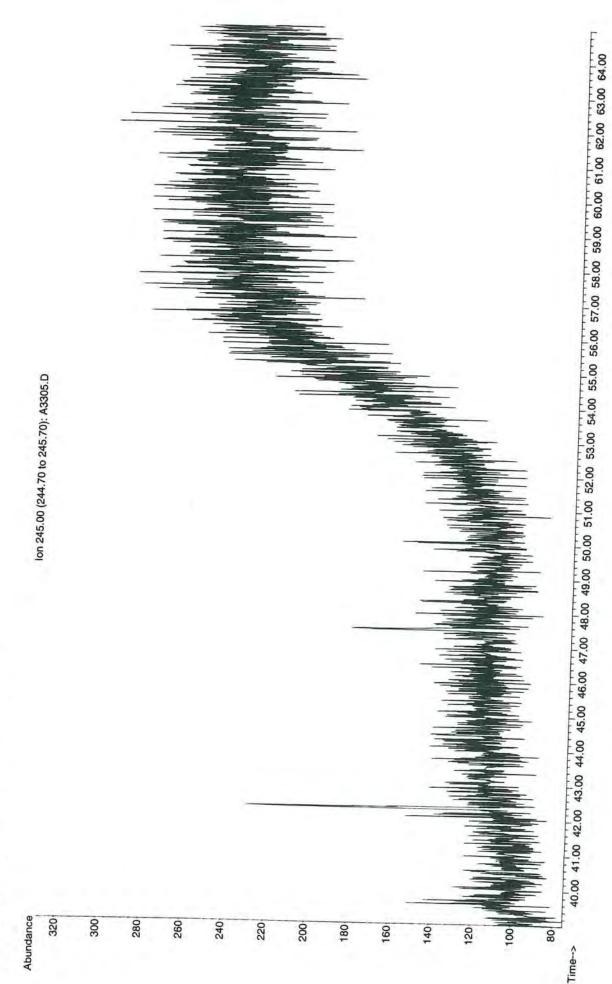


5:15 am using AcqMethod BIOPLUSN.M File : G:\A\DATA\SA0612\A3303.D Operator : DPB Acquired : 24 Oct 2007 5:15 am using Instrument : Inst. A Sample Name: BL043NSC-P(0) Misc Info : North Slope Crude 07-0259 Vial Number: 10



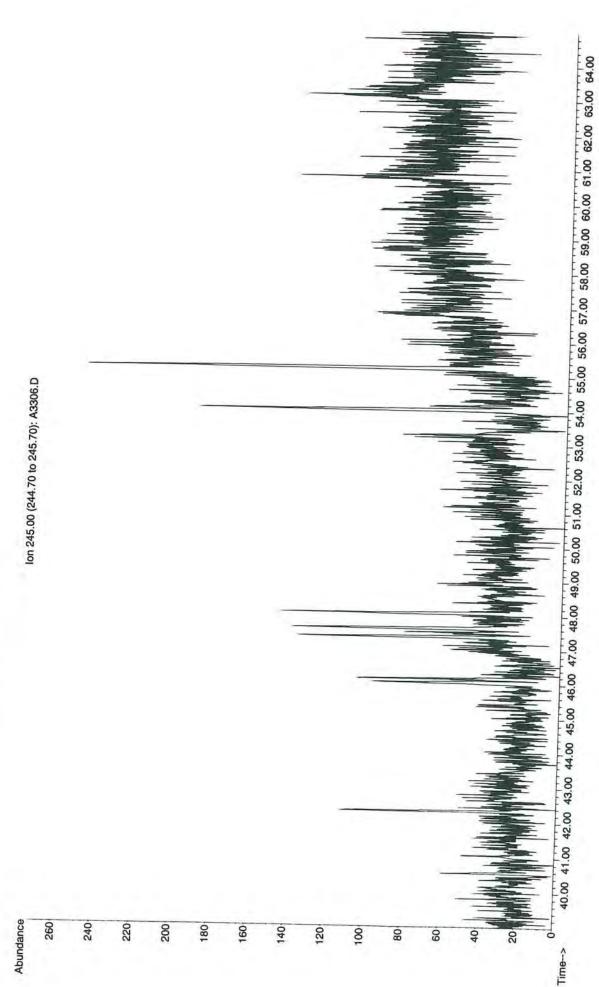


8:13 am using AcqMethod BIOPLUSN : G:\A\DATA\SA0612\A3305.D : DPB : 24 Oct 2007 8:13 am using AcqM : Inst. A : BL033PB-P-MS(5) : Procedural Blank 5-157 07-0259 : 12 Sample Name: Misc Info : Vial Number: Operator Acquired Instrument File





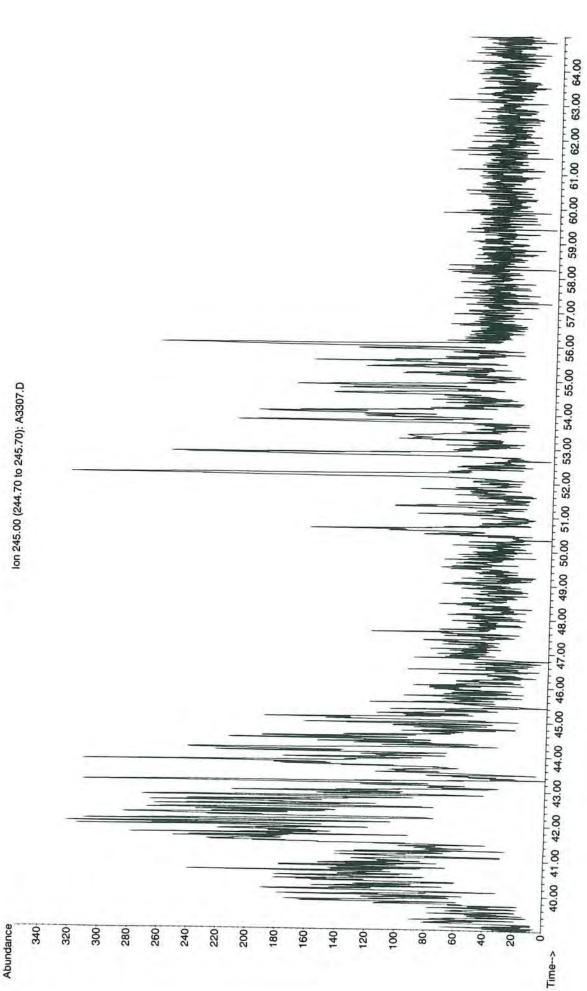
9:42 am using AcqMethod BIOPLUSN.M G:\A\DATA\SA0612\A3306.D DPB 24 Oct 2007 9:42 am using AcqMethod B1 Inst. A BL034LCS-P-MS(5) Laboratory Control Sample 5-157 07-0259 13 Misc Info : Vial Number: Sample Name: Instrument Operator Acquired File





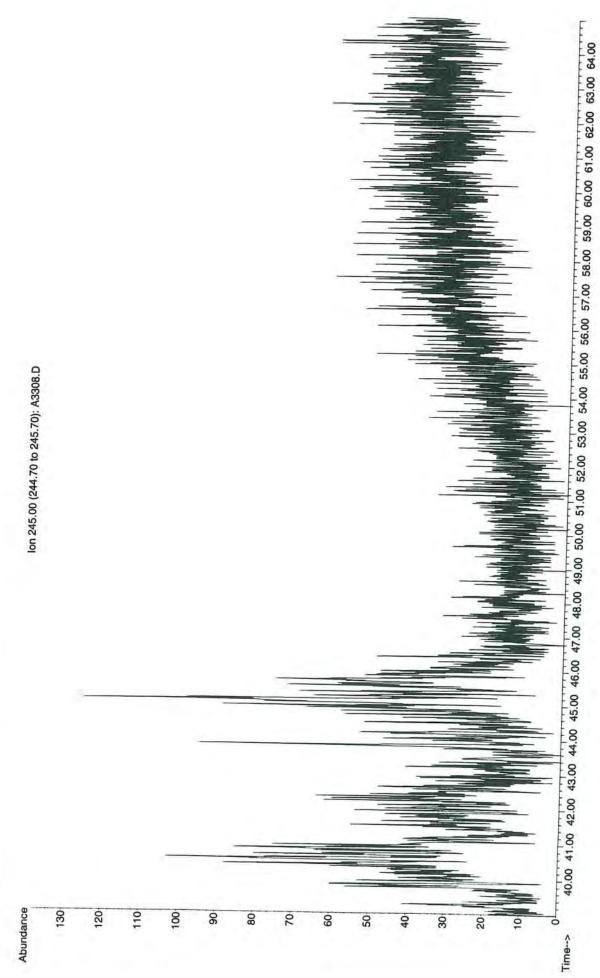


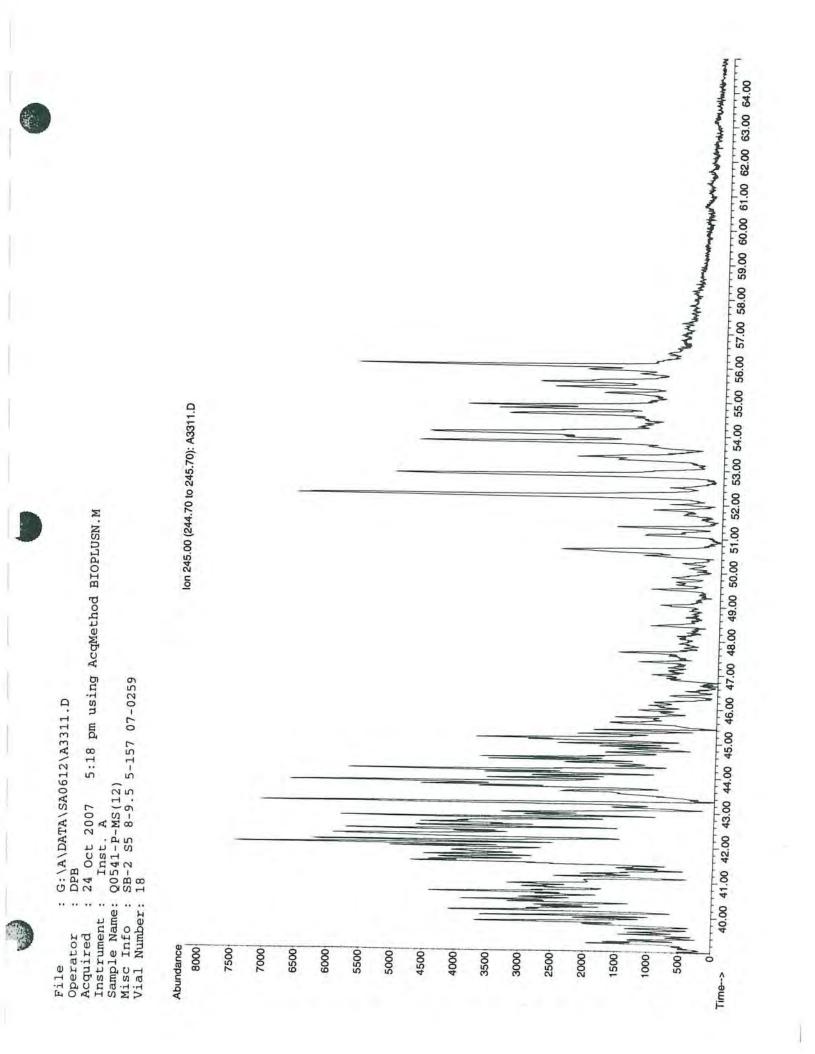
G:\A\DATA\SA0612\A3307.D DPB 24 Oct 2007 11:12 am using AcqMethod BIOPLUSN.M Inst. A Q0541-P-MS-D(13) SB-2 S5 8-9.5 5-157 07-0259 14 Sample Name: (Misc Info : 9 Vial Number: 1 ... •• .. File Operator Acquired Instrument

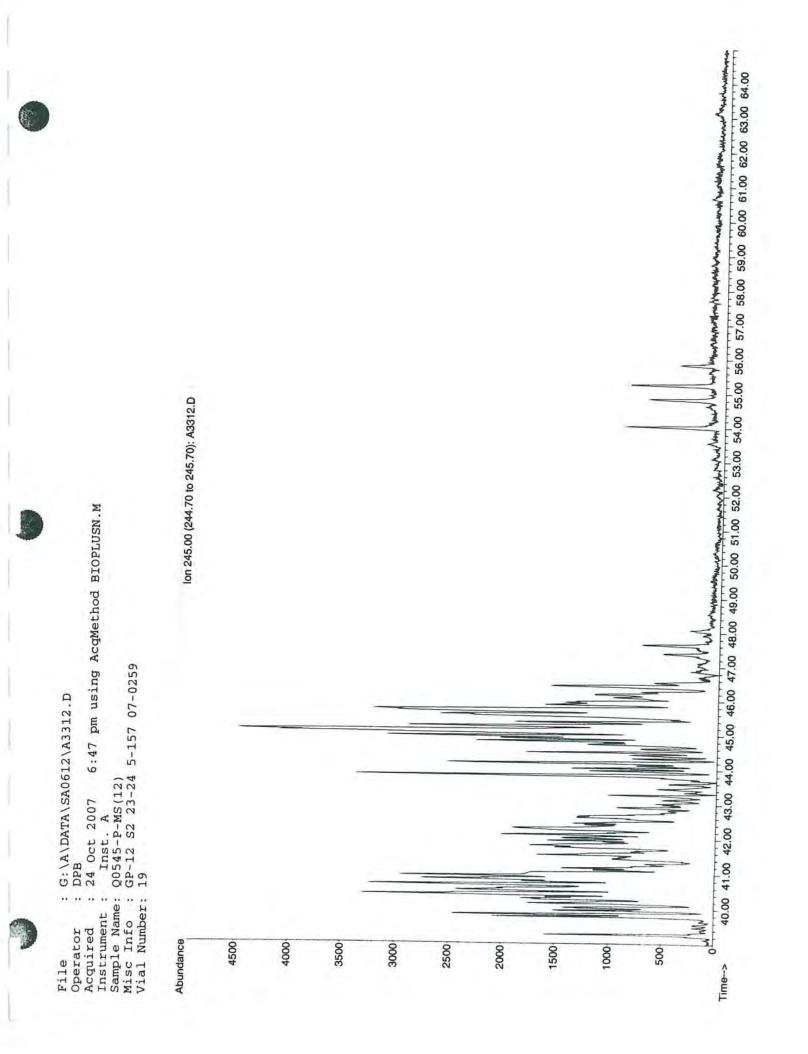


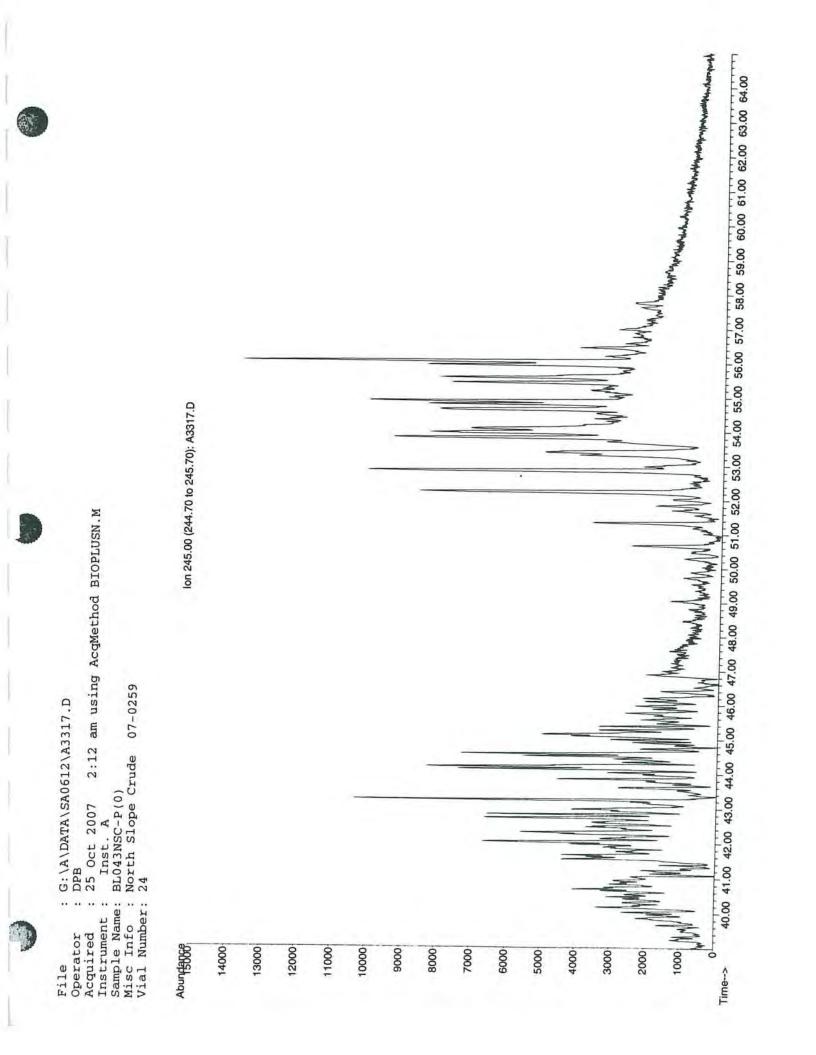


24 Oct 2007 12:41 pm using AcqMethod BIOPLUSN.M Inst. A Q0545-P-MS-D(13) GP-12 S2 23-24 5-157 07-0259 G:\A\DATA\SA0612\A3308.D DPB 15 Sample Name: C Misc Info : C Vial Number: 1 Instrument File Operator Acquired









PAH and Biomarker – SEDIMENT QA/QC SUMMARY Batch 07-0264

PROJECT:	Floyd/Snyder – Gas Works Park
PARAMETER:	Polycyclic Aromatic Hydrocarbons (PAH) and Biomarkers
LABORATORY:	Battelle, Duxbury, MA
MATRIX:	Non-aqueous phase liquid (NAPL)
SAMPLE CUSTODY:	Six soil samples and 1 NAPL were received at Battelle Duxbury Operations (BDO) Laboratory on 10/9/07. Upon Receipt of the samples, the temperature of the cooler was taken and the samples were logged into the laboratory and given unique IDs. The temperature of the cooler upon receipt was slightly below the acceptable range (4°C \pm 2°) at 0.6°C. Also, it was noted that sample Q0540 (MW-9) had some oil residue on the outside of the container as well as the inner side of the bubble wrap. The oil residue was only on the inside of the bubble wrap and did not touch any of the other samples. The client was notified on 10/16/07. The laboratory was instructed to proceed with the analysis. Samples were stored in an access-limited walk-in refrigerator at 4°C until sample preparation could begin.
1	

Reference Method	Method Blank	Surrogate Recovery	LCS Recovery	MS Recovery	MS/MSD Precision	Control Oil % Diff
General NS&T	<5xMDL	40-120% Recovery	40-120% Recovery	40-120% Recovery	≤30% RPD	PD,30% for 90% of the analytes
				MS target spike must be >5x background		
	Method General	MethodBlankGeneral<5xMDL	MethodBlankRecoveryGeneral<5xMDL	MethodBlankRecoveryRecoveryGeneral<5xMDL	Method Blank Recovery Recovery Recovery General <5xMDL	Method Blank Recovery Recovery Recovery Recovery Precision General <5xMDL

METHOD:

The NAPL sample was extracted following general NS&T methods. Approximately 50 mg of oil was weighed and diluted with 10mL of hexane. A portion of the extract was removed and spiked with SIS and IS. One extract was submitted for PAH and petroleum biomarker analysis and the second extract was submitted for SHC and TPH analysis. NAPL sample data is reported on an oil weight basis.

PAH and petroleum biomarkers were measured by gas chromatography –mass spectrometry (GC/MS) in the selected ion mode (SIM). An initial calibration consisting of target analytes was completed prior to analysis to demonstrate the linear range of the analysis. Calibration verification was performed at the beginning and end of each 12 hour period in which samples were analyzed. Concentrations of the PAH and petroleum biomarkers were calculated by the internal standard method. Target PAH were quantified using the average RF generated from the initial calibration. The alkyl homologue PAH series were assigned the RF of the parent PAH, Steranes were assigned the RF of cholestane, and triterpanes were assigned the RF of Moretane.

Note: the reporting limit for the alkyl benzene compounds is orders of magnitude higher than the reporting limits for the rest of the PAH compounds.

HOLDING TIMES: Samples were prepared for analysis in three analytical batches and were extracted within 15 days of sample collection analyzed within 40 days of extraction.

Batch	Extraction Date	Analysis Date		
07-0264	10/18/2007	10/23/2007		

PAH and Biomarker – SEDIMENT QA/QC SUMMARY Batch 07-0264

PROCEDURAL BLANK (PB):	A procedural blank (PB) was prepared with each analytical batch. Blanks were analyzed to ensure the sample extraction and analysis methods were free of contamination.
	07-0264 – No exceedences noted.
	Comments – None.
LABORATORY CONTROL SAMPLE (LCS):	A laboratory control sample (LCS) was prepared with each analytical batch. The percent recoveries of target analytes were calculated to measure data quality in terms of accuracy.
	07-0264 – No exceedences noted.
	Comments – None.
SURROGATE	Five surrogate compounds were added prior to extraction, including naphthalene-d8,
RECOVERY:	acenaphthene-d10, phenanthrene-d10, benzo(a)pyrene-d12, and 5b(H)-chloane. The recovery of the surrogate compound was calculated to measure data quality in terms of accuracy (extraction efficiency).
	07-0264 – Benzo(a)pyrene-d12 is out in NSC, though the NSC still passes for all the analytes and this does not affect the authentic samples.
	Comments - None.
CONTROL OIL:	A control oil (North Slope Crude) was prepared with the analytical batch. The percent difference (PD) between the measured value and the target value was calculated to measure data quality in terms of accuracy.
	07-0264 – No exceedences noted.
	Comments – None.
CALIBRATIONS:	The GC/MS is calibrated with a minimum 5 level curve for all compounds. The percent relative standard deviation (% RSD) between RF for the individual target analytes must be $\leq 25\%$, and the mean RSD of all target analytes must be $\leq 15\%$. Each batch of samples analyzed is bracketed by a continuing calibration verification (CCV) sample, run at a frequency of minimally every 12 hours. The PD between the true value and the CCV should be $\leq 25\%$ for individual analytes. Additionally an initial calibration check (ICC) sample is run immediately after each initial calibration. The PD between the ICC and the initial calibration should be $\leq 25\%$.
	07-0264 – No exceedences noted.
	Comments – None.

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Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	MW-9
Battelle ID	Q0540-P
Sample Type	SA
Collection Date	10/03/07
Extraction Date	10/18/07
Analysis Date	
Analytical Instrument	10/23/07
% Moisture	MS
% Lipid	NA
Matrix	NA
	NAPL
Sample Size	53.70
Size Unit-Basis	MG_OIL
Units	MG/KG_OIL
C3-Alkylbenzenes	957.06
C4-Alkylbenzenes	1229.52
C5-Alkylbenzenes	522.11
C6-Alkylbenzenes	546.55
Benzo(b)thiophene	
C1-benzo(b)thiophenes	230.3
C2-benzo(b)thiophenes	588.57
	770.41
C3-benzo(b)thiophenes	681.89
C4-benzo(b)thiophenes	348.21
Naphthalene	12501.92 D
C1-Naphthalenes	14043.83 D
C2-Naphthalenes	11458.07 D
C3-Naphthalenes	6178.7 D
C4-Naphthalenes	2509.58 D
C1-Biphenyls + Dibenzofuran	1045.39
Biphenyl	705.62
C2-Biphenyls + C1-Dibenzofurans	1651.94
Acenaphthylene	562.91
Acenaphthene	1875.64
Dibenzofuran	251.7
Fluorene	1457.43
C1-Fluorenes	1344.02
C2-Fluorenes	1034.47
C3-Fluorenes	539.52
C4-Fluorenes	485.27
Anthracene	5 B. 17 The 18 1
Phenanthrene	998.56 D
C1-Phenanthrenes/Anthracenes	4785.59 D
C2-Phenanthrenes/Anthracenes	3773.15 D
C3-Phenanthrenes/Anthracenes	1970.53 D
C4-Phenanthrenes/Anthracenes	664.8 D
Retene	208.41 D
	85.3
Dibenzothiophene	359.75
C1-Dibenzothiophenes	547.93
C2-Dibenzothiophenes	508.9
C3-Dibenzothiophenes	278.07
C4-Dibenzothiophenes	97.27
Fluoranthene	979.03
Pyrene	1443.38
C1-Fluoranthenes/Pyrenes	1749.41
C2-Fluoranthenes/Pyrenes	692.53
C3-Fluoranthenes/Pyrenes	264.09
C4-Fluoranthenes/Pyrenes	92.23
C0-Benzo(b)naphthothiophenes	58.98
C1-Benzo(b)naphthothiophenes	
C2-Benzo(b)naphthothiophenes	112.29
C3-Benzo(b)naphthothiophenes	61.47
C1 Paga(b)apphinothiophenes	31.01
C4-Benzo(b)naphthothiophenes	20.71
Benzo(a)anthracene	539.72
Chrysene	538.43

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Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	MW-9
Battelle ID	Q0540-P
Sample Type	SA
Collection Date	10/03/07
Extraction Date	10/18/07
Analysis Date	10/23/07
Analytical Instrument	MS
% Moisture	NA
% Lipid	NA
Matrix	NAPL
Sample Size	53.70
Size Unit-Basis	MG OIL
Units	MG/KG_OIL
C1-Chrysenes	518.15
C2-Chrysenes	226.85
C3-Chrysenes	100.32
C4-Chrysenes	45.7
Benzo(b)fluoranthene	225.58
Benzo(k)fluoranthene	335.92
Benzo(e)pyrene	257.94
Benzo(a)pyrene	512.94
Perylene	85.5
	000.00
Indeno(1,2,3-cd)pyrene	229.39
Indeno(1,2,3-cd)pyrene Dibenz(a,h)anthracene	229.39 56.45

Surrogate Recoveries (%)

Naphthalene-d8	96
Acenaphthene-d10	90
Phenanthrene-d10	86
Benzo(a)pyrene-d12	120
5b(H)-Cholane	115

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Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	Procedural Blank	
Battelle ID	BL057PB-P	
Sample Type	PB	
Collection Date	10/18/07	
Extraction Date	10/18/07	
Analysis Date	10/23/07	
Analytical Instrument	MS	
% Moisture	NA	
% Lipid		
Matrix	NA	
Sample Size	OIL	
	50.00	
Size Unit-Basis Units	MG_OIL	
Units	MG/KG_OIL	
C3-Alkylbenzenes	U	
C4-Alkylbenzenes	U	
C5-Alkylbenzenes	U	
C6-Alkylbenzenes	Ŭ	
Benzo(b)thiophene	Ŭ	
C1-benzo(b)thiophenes		
C2-benzo(b)thiophenes	U	
	0	
C3-benzo(b)thiophenes	U	
C4-benzo(b)lhiophenes	U	
Naphthalene	0.09 J	
C1-Naphthalenes	U	
C2-Naphthalenes	Ŭ	
C3-Naphthalenes	Ŭ	
C4-Naphthalenes	Ŭ	
C1-Biphenyls + Dibenzofuran	U	
Biphenyl	U	
C2-Biphenyls + C1-Dibenzofurans		
Acenaphthylene	u	
	U	
Acenaphthene	U	
Dibenzofuran	U	
Fluorene	U	
C1-Fluorenes	Ŭ	
C2-Fluorenes	U	
C3-Fluorenes	Ũ	
C4-Fluorenes	Ŭ	
Anthracene	ŭ	
Phenanthrene		
C1-Phenanthrenes/Anthracenes	U	
C2-Phenanthrenes/Anthracenes	U	
C2-Frienanthrenes/Anthracenes	U	
C3-Phenanthrenes/Anthracenes	U	
C4-Phenanthrenes/Anthracenes	U	
Retene	U	
Dibenzothiophene	Ū	
C1-Dibenzothiophenes	Ŭ	
C2-Dibenzothiophenes	Ŭ	
C3-Dibenzothiophenes		
C4-Dibenzothiophenes	U	
	υ	
Fluoranthene	U	
Pyrene	Ŭ	
C1-Fluoranthenes/Pyrenes	U	
C2-Fluoranthenes/Pyrenes	Ŭ	
C3-Fluoranthenes/Pyrenes	ũ	
C4-Fluoranthenes/Pyrenes	ŭ	
CO-Benzo(b)naphthothiophenes	Ŭ	
C1-Benzo(b)naphthothiophenes		
2-Benzo(b)naphthothiophenes	U	
3-Benzo(b)naphthothicshores	U	
3-Benzo(b)naphthothiophenes	U	
24-Benzo(b)naphthothiophenes	U	
lenzo(a)anthracene	U	
Chrysene	Ŭ	

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Client ID	Procedural Blank		
Battelle ID	BL057PB-P		
Sample Type	PB		
Collection Date	10/18/07		
Extraction Date	10/18/07		
Analysis Date	10/23/07		
Analytical Instrument	MS		
% Moisture	NA		
% Lipid	NA		
Matrix	OIL		
Sample Size	50.00		
Size Unit-Basis	MG_OIL		
Units	MG/KG_OIL		
C1-Chrysenes	U		
C2-Chrysenes	Ŭ		
C3-Chrysenes	ŭ		
C4-Chrysenes	Ŭ		
Benzo(b)fluoranthene	U U		
Benzo(k)fluoranthene	Ŭ		
Benzo(e)pyrene	Ŭ		
Benzo(a)pyrene	Ŭ		
Perylene	Ŭ		
Indeno(1,2,3-cd)pyrene	Ŭ		
Dibenz(a,h)anthracene	Ŭ		
Benzo(g,h,i)perylene	U		
Total PAH	0.09 J		
	0.09 0		
Surrogate Recoveries (%)			
Naphthalene-d8	106		
Acenaphthene-d10	97		
Phenanthrene-d10	96		
Benzo(a)pyrene-d12	109		
5b(H)-Cholane	95		
and the second sec	55		

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Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	Laboratory Contro Sample					
Battelle ID	BL058LCS-F					
Sample Type	LCS					
Collection Date	10/18/07					
Extraction Date	10/18/07					
Analysis Date	10/23/07					
Analytical Instrument	MS					
% Moisture	NA					
% Lipid	NA					
Matrix	OIL					
Sample Size	NA					
Size Unit-Basis	NA					
Units	NG		Target	% Recovery	Qualifier	
C3-Alkylbenzenes		14				
C4-Alkylbenzenes		U				
C5-Alkylbenzenes		U				
C6-Alkylbenzenes		UU				
Benzo(b)thiophene	1697.65	0	0000.00			
C1-benzo(b)thiophenes	1097.05	- n	2006.80	85		
C2-benzo(b)thiophenes		U				
C3-benzo(b)thiophenes		UU				
C4-benzo(b)thiophenes		ŭ				
Naphthalene	1948.35	U	2000 10	07		
C1-Naphthalenes	1940.35	υ	2000.40	97		
C2-Naphthalenes		U				
C3-Naphthalenes		U				
C4-Naphthalenes		Ŭ				
C1-Biphenyls + Dibenzofuran		ŭ				
Biphenyl	1681.44	0	2003.40	84		
C2-Biphenyls + C1-Dibenzofurans	1001.44	U	2003.40	04		
Acenaphthylene	1859.47	U	2001.80	93		
Acenaphthene	1962.93		2001.30	98		
Dibenzofuran	1739.28		2003.80	87		
Fluorene	1864.93		2001.10	93		
C1-Fluorenes	1.	U	2001.10	55		
C2-Fluorenes		Ŭ				
C3-Fluorenes		Ũ				
C4-Fluorenes		U				
Anthracene	1779.37		2000.30	89		
Phenanthrene	1781.84		2001.00	89		
C1-Phenanthrenes/Anthracenes		U		105		
C2-Phenanthrenes/Anthracenes		U				
C3-Phenanthrenes/Anthracenes		U				
C4-Phenanthrenes/Anthracenes		U				
Retene		U				
Dibenzothiophene	1570.79		2002.00	78		
C1-Dibenzothiophenes		U				
C2-Dibenzothiophenes		U				
C3-Dibenzothiophenes		U				
C4-Dibenzothiophenes	- Alternation	U				
Fluoranthene Pyrene	1732.39		2001.00	87		
C1-Fluoranthenes/Pyrenes	1782.5		2000.70	89		
		U				
C2-Fluoranthenes/Pyrenes C3-Fluoranthenes/Pyrenes		U				
C4-Fluoranthenes/Pyrenes		U				
C0-Benzo(b)naphthothiophenes		U				
C1-Benzo(b)naphthothiophenes		U				
C2-Benzo(b)naphthothiophenes		U				
C3-Benzo(b)naphthothiophenes		U				
C4-Benzo(b)naphthothiophenes		U				
Benzo(a)anthracene	1698.46	U	2000 50	05		
	1090.40		2000.50	85		

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Client ID	Laboratory Control Sample					
Detter ID						
Battelle ID	BL058LCS-P					
Sample Type	LCS					
Collection Date	10/18/07					
Extraction Date	10/18/07					
Analysis Date	10/23/07					
Analytical Instrument	MS					
% Moisture	NA					
% Lipid	NA					
Matrix	OIL					
Sample Size	NA					
Size Unit-Basis	NA.					
Units	NG	-	Target %	& Recovery	Qualifier	
Chrysene	1712.45		2000.80	86		
C1-Chrysenes	.,	U	2000.00	00		
C2-Chrysenes		Ŭ				
C3-Chrysenes		ŭ				
C4-Chrysenes		ŭ				
Benzo(b)fluoranthene	1859.09	0	2001.70	93		
Benzo(k)/luoranthene	2011.19		2001.10	101		
Benzo(e)pyrene	1721.05		2002.80	86		
Benzo(a)pyrene	1965.52		2001.60	98		
Perylene	1873.42		2004.10	93		
ndeno(1,2,3-cd)pyrene	1851.82		2001.00	93		
Dibenz(a,h)anthracene	1779.91		2001.10	89		
Benzo(g,h,i)perylene	1811.31		2000.60	91		
Total PAH	37987.51		2000.00	91		
Surrogate Recoveries (%)						
laphthalene-d8	104					
cenaphthene-d10	96					
henanthrene-d10	95					
enzo(a)pyrene-d12	110					
	110					

Battelle The Business of Innovation

Client ID	GO98: North Slope Crude			
Battelle ID	BL060NSC-P			
Sample Type	NSC			
Collection Date	10/18/07			
Extraction Date	10/18/07			
Analysis Date	10/23/07			
Analytical Instrument	MS			
% Moisture	NA			
% Lipid	NA			
Matrix	OIL			
Sample Size	5.04			
Size Unit-Basis	MG_OIL			
Units	MG/KG_OIL	Target % I	Difference	Qualifier
C3-Alkylbenzenes	1560.8			
C4-Alkylbenzenes	1216.22			
C5-Alkylbenzenes	685.39			
C6-Alkylbenzenes	430.56			
Benzo(b)thiophene	12.17			
C1-benzo(b)thiophenes	39.72			
C2-benzo(b)thiophenes	72.43	95,74	04.0	
C3-benzo(b)thiophenes	127.49	132.67	24.3 3.9	
C4-benzo(b)thiophenes	85.27	96.72		
Naphthalene	689.11	740.29	11.8	
C1-Naphthalenes	1395.18	1516.04	6.9	
C2-Naphthalenes	1805.37	2000.10	8.0 9.7	
C3-Naphthalenes	1399.36	1526.96	8.4	
C4-Naphthalenes	755.68	898.03	15.9	
C1-Biphenyls + Dibenzofuran	318.25	030.03	15.9	
Biphenyl	204.89	220.82	70	
C2-Biphenyls + C1-Dibenzofurans	473.11	220.02	7.2	
Acenaphthylene	473.11 U			
Acenaphthene	12.4	14.50	14.5	
Dibenzofuran	64.18	77.75		
Fluorene	95.2	92.51	17.5	
C1-Fluorenes	200.32	227.01	11.8	
C2-Fluorenes	287.55	367.09	21.7	
C3-Fluorenes	250.22	326.32	23.3	
C4-Fluorenes	157.75	020.02	20.0	
Anthracene	U			
Phenanthrene	228.28	249.49	8.5	
C1-Phenanthrenes/Anthracenes	497.37	549.17	9.4	
C2-Phenanthrenes/Anthracenes	574.96	642.72	10.5	
C3-Phenanthrenes/Anthracenes	390.44	446.11	12.5	
C4-Phenanthrenes/Anthracenes	142.25	180.02	21.0	
Retene	43.86	100.02	11.0	
Dibenzothiophene	194.31	210.35	7.6	
C1-Dibenzothiophenes	362.86	409.03	11.3	
C2-Dibenzothiophenes	498.7	551.46	9.6	
C3-Dibenzothiophenes	408.82	471.36	13.3	
C4-Dibenzothiophenes	221.26	243.11	9.0	
Fluoranthene	3.55	2.131.1		
Pyrene	15.21	12.99	17.1	
C1-Fluoranthenes/Pyrenes	61.96	70.92	12.6	
C2-Fluoranthenes/Pyrenes	119.12	117.89	1.0	
C3-Fluoranthenes/Pyrenes	141.73	137.25	3.3	
C4-Fluoranthenes/Pyrenes	98.19		0.0	
C0-Benzo(b)naphthothiophenes	39.22			
C1-Benzo(b)naphthothiophenes	153.8			
C2-Benzo(b)naphthothiophenes	189.65			
C3-Benzo(b)naphthothiophenes	128.56			
C4-Benzo(b)naphthothiophenes	78.36			
Benzo(a)anthracene	4.17			
	Vi-			

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Client ID	GO98: North Slope Crude					
Battelle ID	BL060NSC-P					
Sample Type	NSC					
Collection Date	10/18/07					
Extraction Date	10/18/07					
Analysis Date	10/23/07					
Analytical Instrument	MS					
% Moisture	NA					
% Lipid	NA					
Matrix	OIL					
Sample Size	5.04					
Size Unit-Basis	MG_OIL					
Units	MG/KG_OIL		Target % [Difference	Qualifier	
Chrysene	47.25		47.18			
C1-Chrysenes	47.25		78.82	0,1		
C2-Chrysenes	102.3		102.67	0.2		
C3-Chrysenes	88.83		85.36	4.1		
C4-Chrysenes	55.14		61.99	4.1		
Benzo(b)fluoranthene	5.89		6.08	3.1		
Benzo(k)fluoranthene	0.05	U	0.00	5.1		
Benzo(e)pyrene	13.72	0	12.88	6.5		
Benzo(a)pyrene	1.09	J	12.00	0,5		
Perylene	1.00	Ŭ				
Indeno(1,2,3-cd)pyrene		ŭ				
Dibenz(a,h)anthracene	1.09	J				
Benzo(g,h,i)perylene	3.61		3.44	4.9		
Total PAH	11422		5,11	4.5		
Surrogate Recoveries (%)						
Naphthalene-d8	104					
Acenaphthene-d10	92					
Phenanthrene-d10	90					
		N				
Benzo(a)pyrene-d12 5b(H)-Cholane	126 96	N				

Procedural Blank (BL057PB-P)

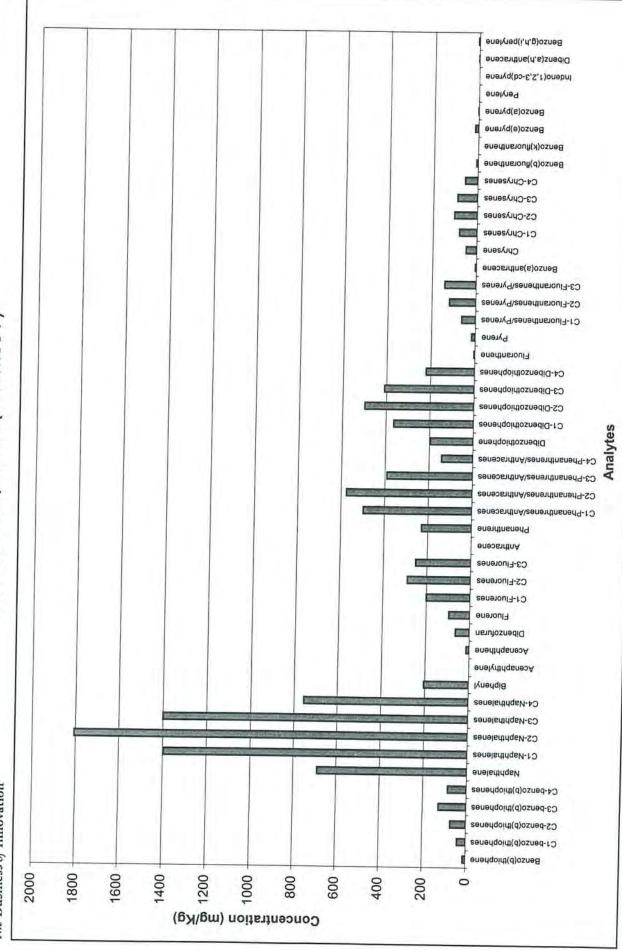
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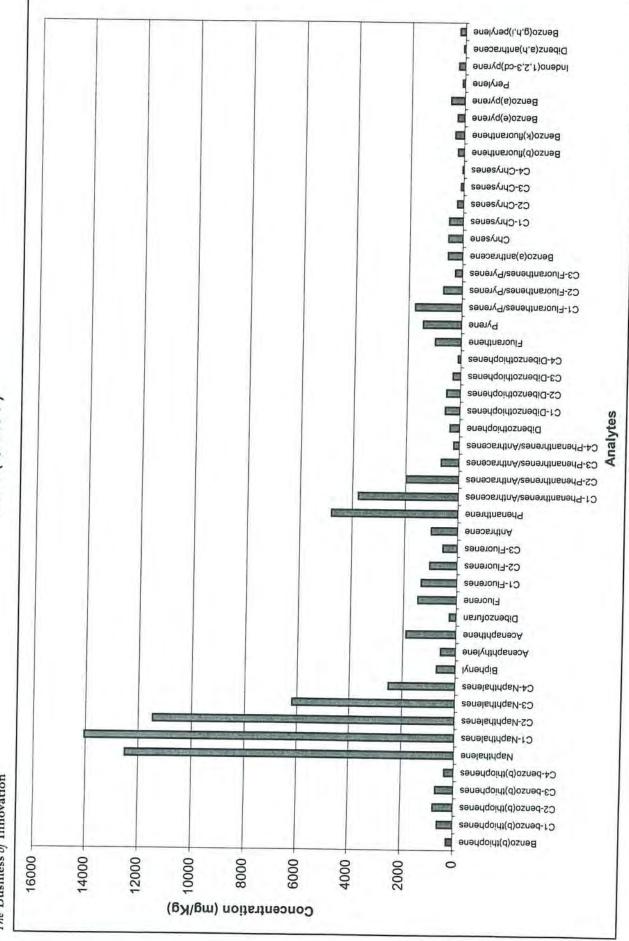


GO98: North Slope Crude (BL060NSC-P)





MW-9 (Q0540-P)



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Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	MW-9
Battelle ID	Q0540-P
Sample Type	SA
Collection Date	10/03/07
Extraction Date	10/18/07
Analysis Date	
Analytical Instrument	10/23/07
% Moisture	MS
% Lipid	NA
Matrix	NA
Sample Size	NAPL
Size Unit-Basis	53.70
Units	MG_OIL
Onits	MG/KG_OIL
C23 Tricyclic Terpane	44.57
C29 Tricyclic Terpane -22S	13.91
C29 Tricyclic Terpane -22R	12.55
18a(H)-22,29,30-Trisnomeohopane -TS	8.56
17a(H)-22,29,30-Trisnorhopane -TM	12.67
30-Norhopane	35.59
18a(H) & 18b(H)-Oleananes	13.97
Hopane	65.08
30-Homohopane -22S	23.21
30-Homohopane -22R	13.35
13b(H),17a(H)-20S-Diacholestane	
13b(H),17a(H)-20R-Diacholestane	78.45
14a(H),17a(H)-20R-methylcholestane	45.43
14a(H),17a(H)-20S-Ethylcholestane	102.96
14a(H),17a(H)-20R-Ethylcholestane	40.52
C21-TAS	59.44
C26-TAS(20S)	10.91
C26,C27-TAS	18.2
C27-TAS(20R)	58.72
C28-TAS(20S)	30.58
C28-TAS(20R)	18.79
C21-MAS	14.16
C22-MAS	7.93
C27-MAS	3.89
C27-20R-MAS	3.14
	18.71
C27-20S-MAS	14.78
C28-20S-MAS	32.2
C27-C2920S/R-MAS	36.37
C29-20S-MAS	33.57
C29-20R-MAS	24.77
TAS_245	υ
MAS 239	U

Surrogate Recoveries (%)

Naphthalene-d8	96
Acenaphthene-d10	90
Phenanthrene-d10	86
Benzo(a)pyrene-d12	120
5b(H)-Cholane	115

The Business of Innovation

Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

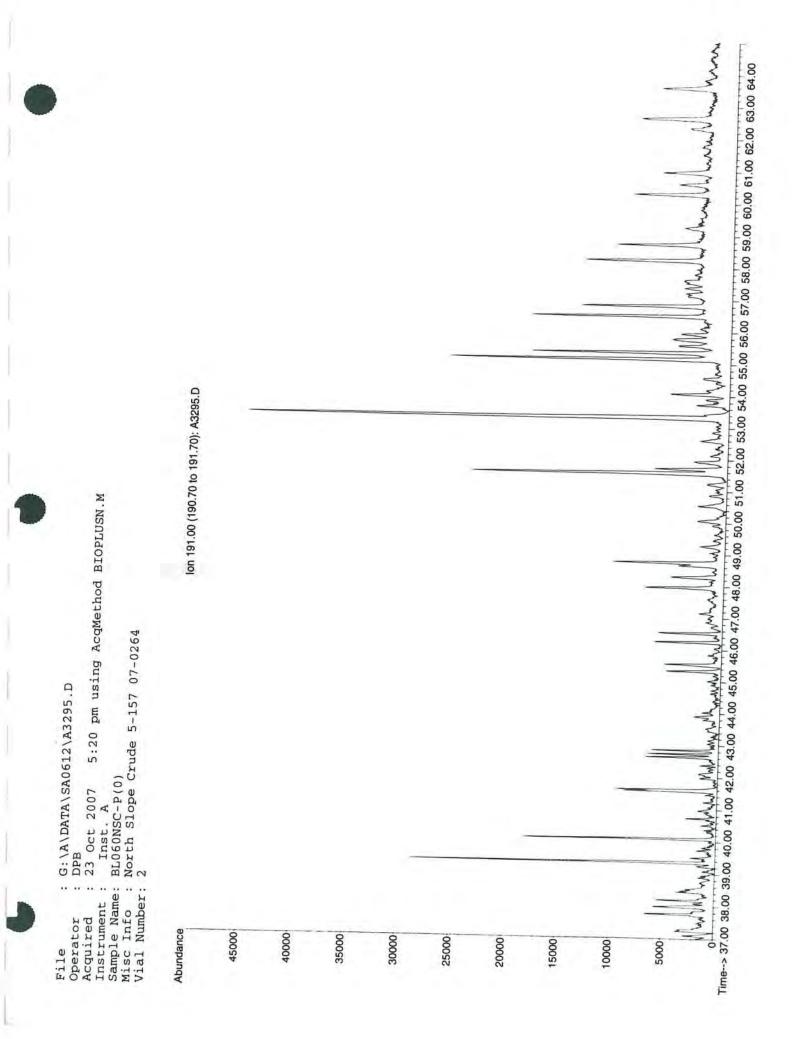
Client ID	Procedural Blank	
B-#-#- ID	100 AT 11 AT 10	
Battelle ID	BL057PB-P	
Sample Type	PB	
Collection Date	10/18/07	
Extraction Date	10/18/07	
Analysis Date	10/23/07	
Analytical Instrument	MS	
% Moisture	NA	
% Lipid	NA	
Matrix	OIL	
Sample Size		
Size Unit-Basis	50.00	
Units	MG_OIL	
Units	MG/KG_OIL	
C23 Tricyclic Terpane	υ	
C29 Tricyclic Terpane -22S	U	
C29 Tricyclic Terpane -22R		
	U	
18a(H)-22,29,30-Trisnomeohopane -TS	U	
17a(H)-22,29,30-Trisnorhopane -TM	U	
30-Norhopane	U	
18a(H) & 18b(H)-Oleananes	Ŭ	
Hopane	U	
30-Homohopane -22S	U	
30-Homohopane -22R	Ū	
13b(H),17a(H)-20S-Diacholestane	ŭ	
13b(H),17a(H)-20R-Diacholestane	ŭ	
14a(H),17a(H)-20R-methylcholestane	Ŭ	
14a(H),17a(H)-20S-Ethylcholestane		
14a(H),17a(H)-20R-Ethylcholestane	U	
C21-TAS	U	
	U	
C26-TAS(20S)	U	
C26,C27-TAS	U	
C27-TAS(20R)	U	
C28-TAS(20S)	U	
C28-TAS(20R)	U	
C21-MAS	U	
C22-MAS	Ŭ	
C27-MAS	Ũ	
C27-20R-MAS	ŭ	
C27-20S-MAS	Ŭ	
C28-20S-MAS	Ŭ	
C27-C2920S/R-MAS	0	
C29-20S-MAS	U	
C29-20R-MAS	U	
	U	
TAS_245	U	
MAS_239	U	
Surrogate Recoveries (%)		
Naphthalene-d8	100	
Acenaphthene-d10	106	
	97	
Phenanthrene-d10	96	
Benzo(a)pyrene-d12	109	
5b(H)-Cholane	95	

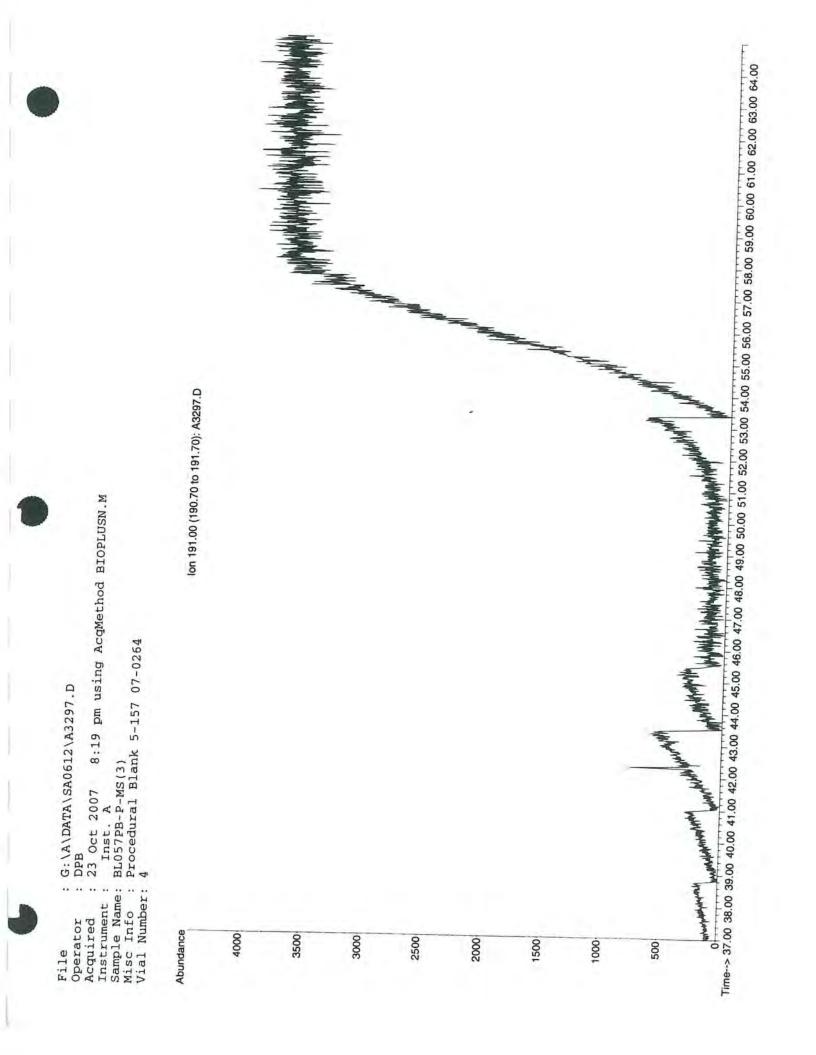
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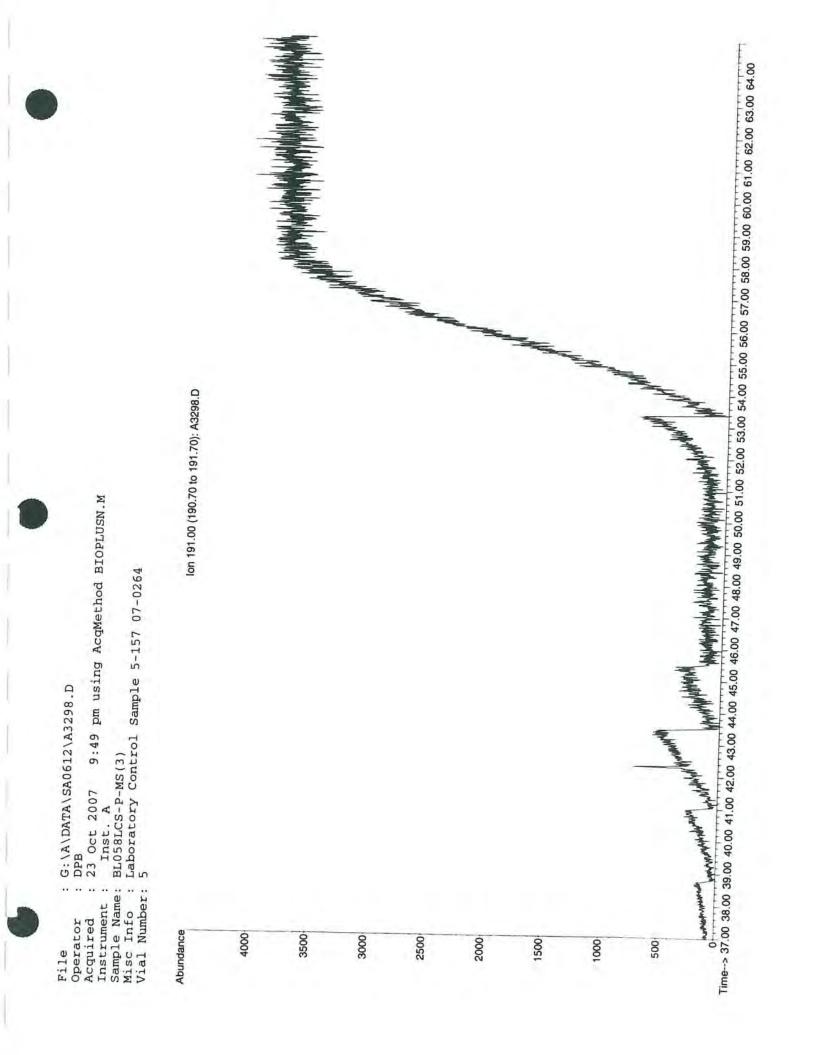
Client ID	Laboratory Control Sample			
	Sample			
Battelle ID	BL058LCS-P			
Sample Type	LCS			
Collection Date	10/18/07			
Extraction Date	10/18/07			
Analysis Date	10/23/07			
Analytical Instrument	MS			
% Moisture	NA			
% Lipid	NA			
Matrix	OIL			
Sample Size	NA			
Size Unit-Basis	NA			
Units	NG	Target % Recovery	Qualifier	
Call States and the states of		raiget in Recovery	Quanter	
C23 Tricyclic Terpane	U			
C29 Tricyclic Terpane -22S	U			
C29 Tricyclic Terpane -22R	U			
18a(H)-22,29,30-Trisnomeohopane -TS	U			
17a(H)-22,29,30-Trisnorhopane -TM	U			
30-Norhopane	U			
18a(H) & 18b(H)-Oleananes	U			
Hopane	U			
30-Homohopane -22S	Ũ			
30-Homohopane -22R	Ū			
13b(H),17a(H)-20S-Diacholestane	Ũ			
13b(H),17a(H)-20R-Diacholestane	Ũ			
14a(H),17a(H)-20R-methylcholestane	Ũ			
14a(H),17a(H)-20S-Ethylcholestane	Ũ			
14a(H),17a(H)-20R-Ethylcholestane	Ŭ			
C21-TAS	Ŭ			
C26-TAS(20S)	Ŭ			
C26,C27-TAS	Ŭ			
C27-TAS(20R)	ŭ			
C28-TAS(20S)	Ŭ			
C28-TAS(20R)	Ŭ			
C21-MAS	Ŭ			
C22-MAS	ŭ			
C27-MAS	ŭ			
C27-20R-MAS	Ŭ			
C27-20S-MAS	Ŭ			
C28-20S-MAS	Ŭ			
C27-C2920S/R-MAS	Ŭ			
C29-20S-MAS	Ŭ			
C29-20R-MAS	Ŭ			
TAS 245	Ŭ			
MAS_239	Ŭ			
	U			
Surrogate Recoveries (%)				
Naphthalene-d8	104			
Acenaphthene-d10	96			
Phenanthrene-d10	95			
Benzo(a)pyrene-d12	110			
5b(H)-Cholane	88			

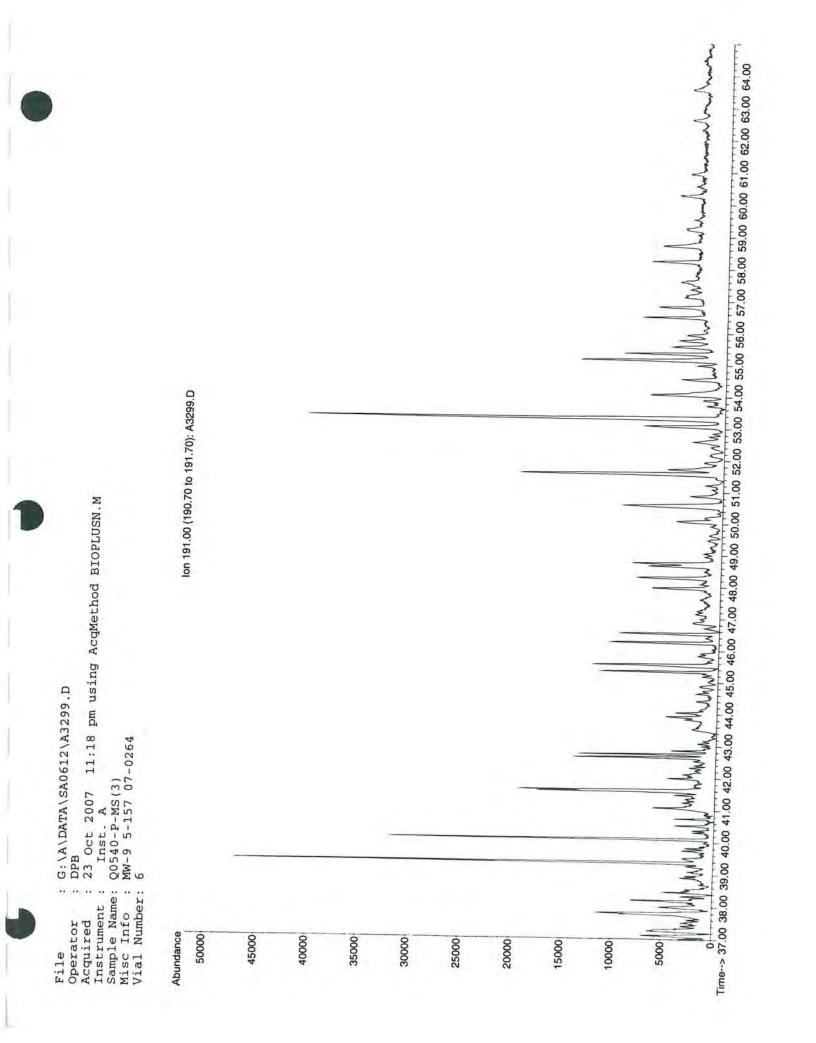
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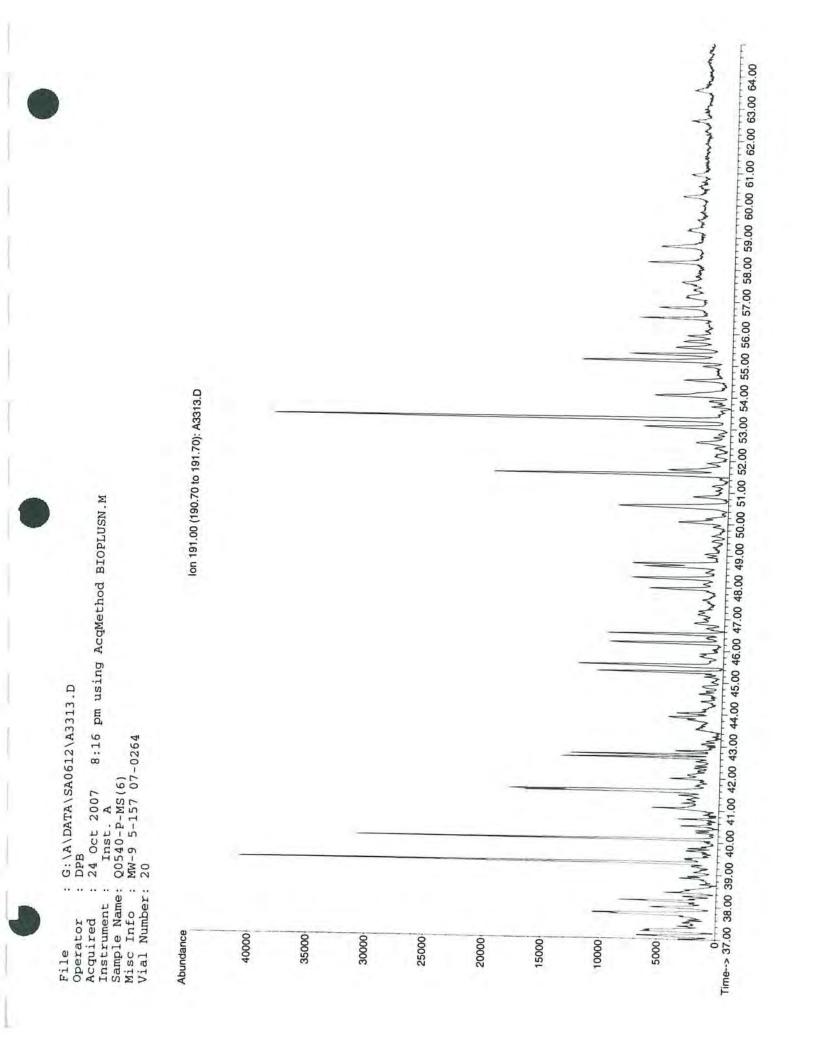
	GO98: North Slope					
Client ID	Crude					
Battelle ID	BL060NSC-P					
Sample Type						
Collection Date	NSC					
Extraction Date	10/18/07					
	10/18/07					
Analysis Date	10/23/07					
Analytical Instrument	MS					
% Moisture	NA					
% Lipid	NA					
Matrix	OIL					
Sample Size	5.04					
Size Unit-Basis	MG OIL					
Units	MG/KG_OIL	1	Target %	Difference	Qualifier	
C23 Tricyclic Terpane	00.50		(in 15)	6.4.5		
C29 Tricyclic Terpane -22S	39.58		47.76	17.1		
	11.85		14.70	19.4		
C29 Tricyclic Terpane -22R	11.1		14.64	24.2		
18a(H)-22,29,30-Trisnomeohopane -TS	15.34		15.96	3.9		
17a(H)-22,29,30-Trisnorhopane -TM	22.14		24.82	10.8		
30-Norhopane	61.9		69.58	11.0		
18a(H) & 18b(H)-Oleananes	0.88	J				
Hopane	106.92		120.14	11.0		
30-Homohopane -22S	55.79		59.93	6.9		
30-Homohopane -22R	39.79		39.69	0.3		
13b(H),17a(H)-20S-Diacholestane	38.7		44.18	12.4		
13b(H),17a(H)-20R-Diacholestane	21.96		25.52	13.9		
14a(H),17a(H)-20R-methylcholestane	28.93		33.94	14.8		
14a(H),17a(H)-20S-Ethylcholestane	33.42		35.93	7.0		
14a(H),17a(H)-20R-Ethylcholestane	36.01		39.17	8.1		
C21-TAS	17.14		00.17	0.1		
C26-TAS(20S)	12.87					
C26,C27-TAS	52.98					
C27-TAS(20R)	31.83					
C28-TAS(20S)	28.99					
C28-TAS(20R)	26.51					
C21-MAS	5.01					
C22-MAS	3.29					
C27-MAS						
C27-20R-MAS	4.68					
C27-20S-MAS	7.65					
C28-20S-MAS	1.36					
C27-C2920S/R-MAS	14.61					
C29-20S-MAS	13.91					
C29-20R-MAS	3.56					
	9.39					
TAS_245	U					
MAS_239	U					
Surrogate Recoveries (%)						
Nachthalana da						
Naphthalene-d8	104					
Acenaphthene-d10	92					
Phenanthrene-d10	90					
Benzo(a)pyrene-d12	126 N					
5b(H)-Cholane	96					

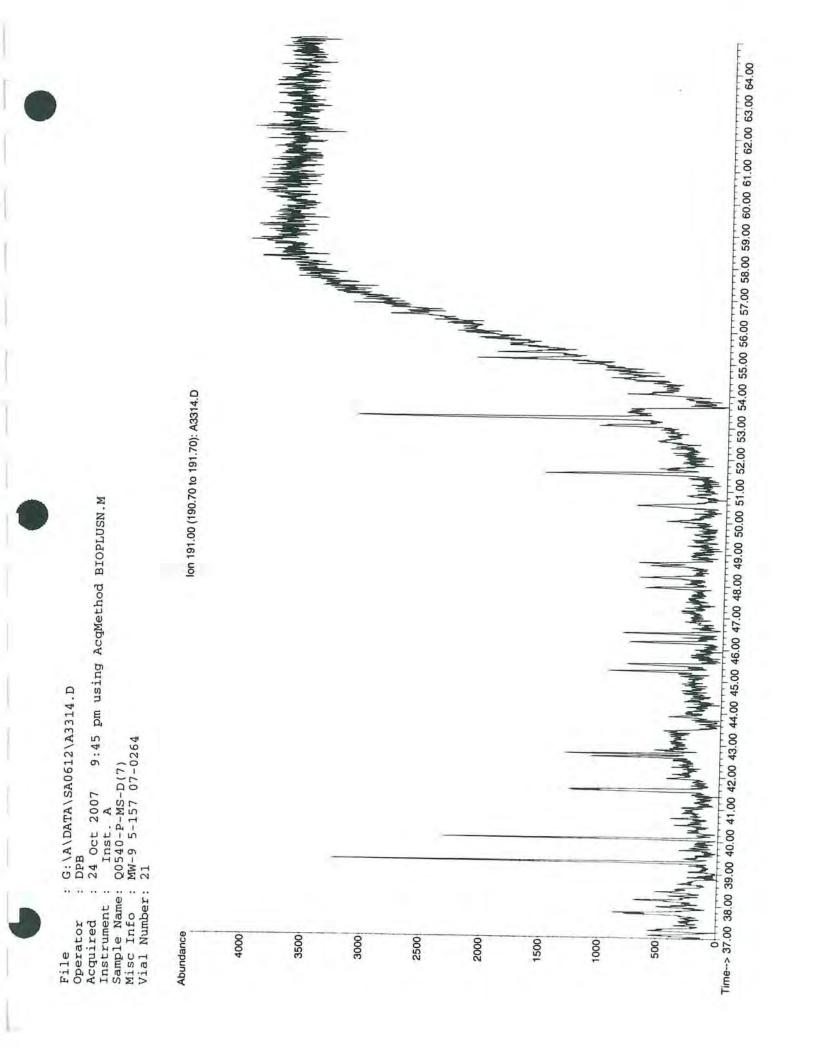


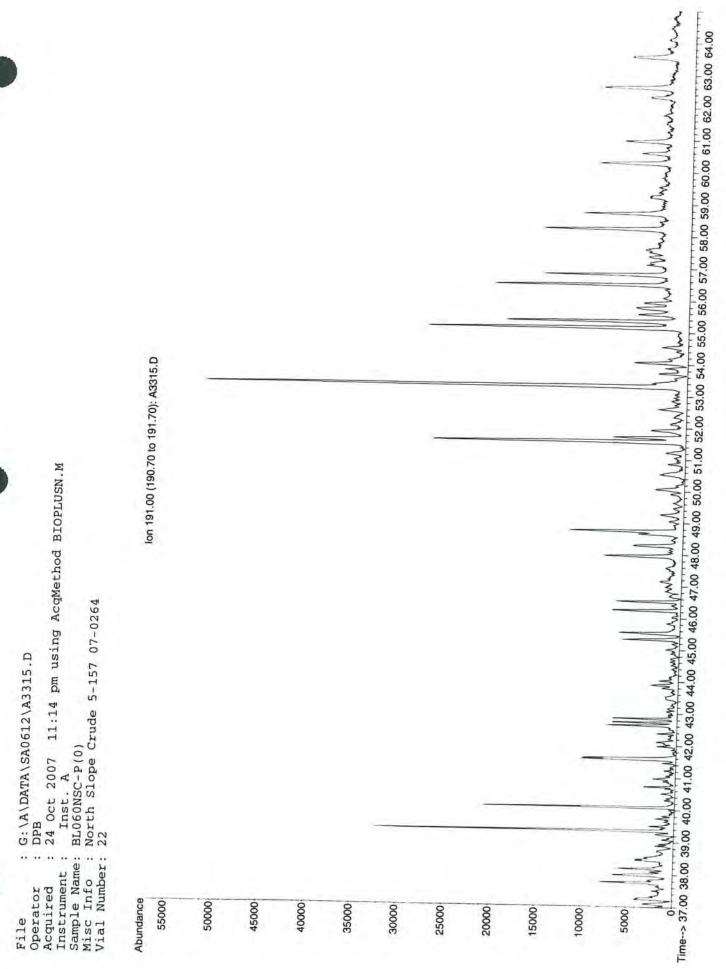


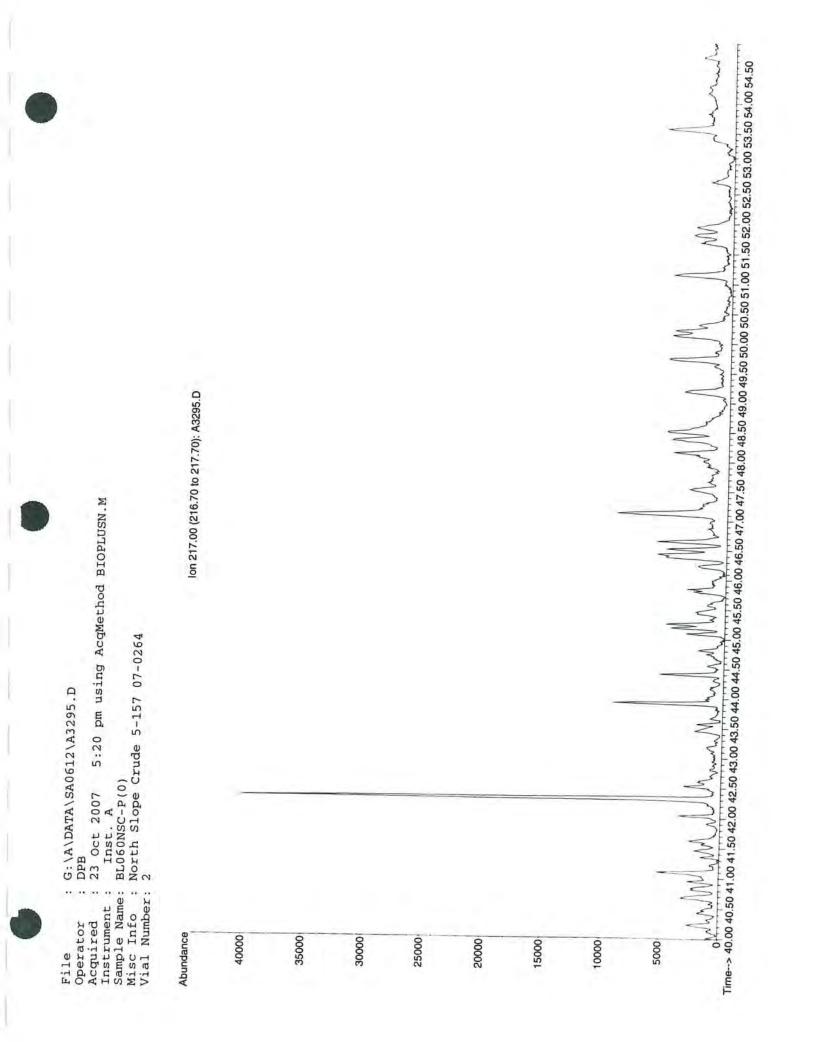










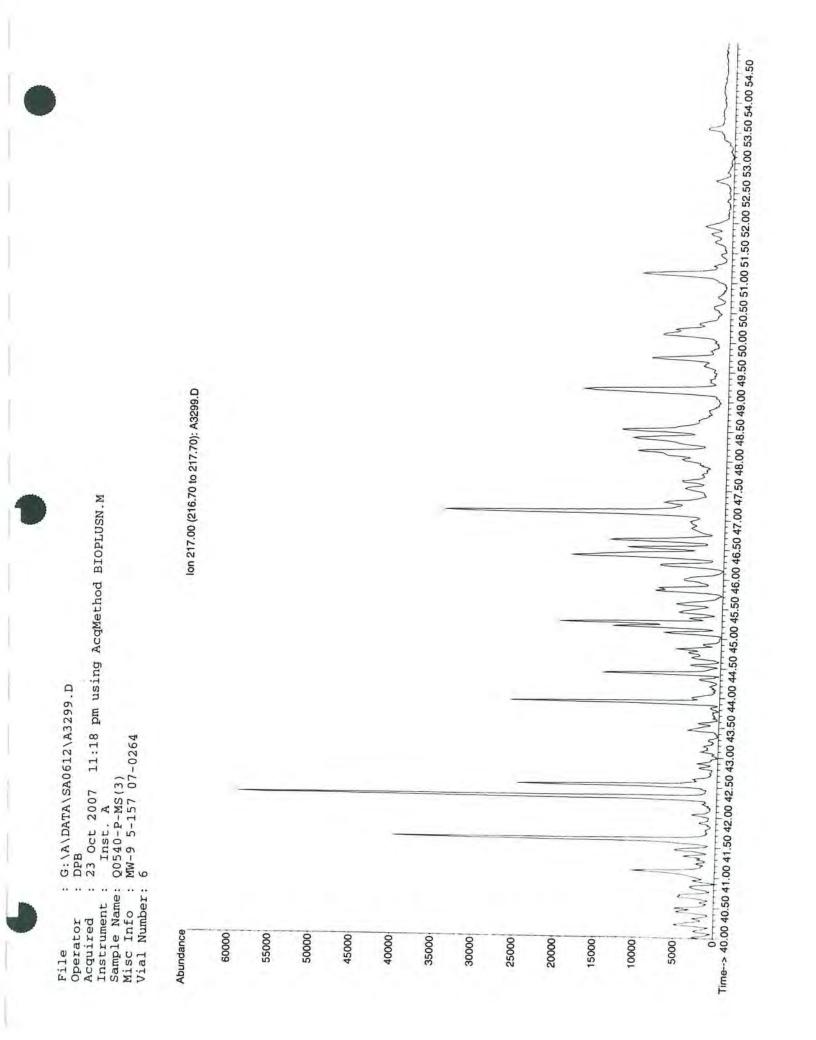


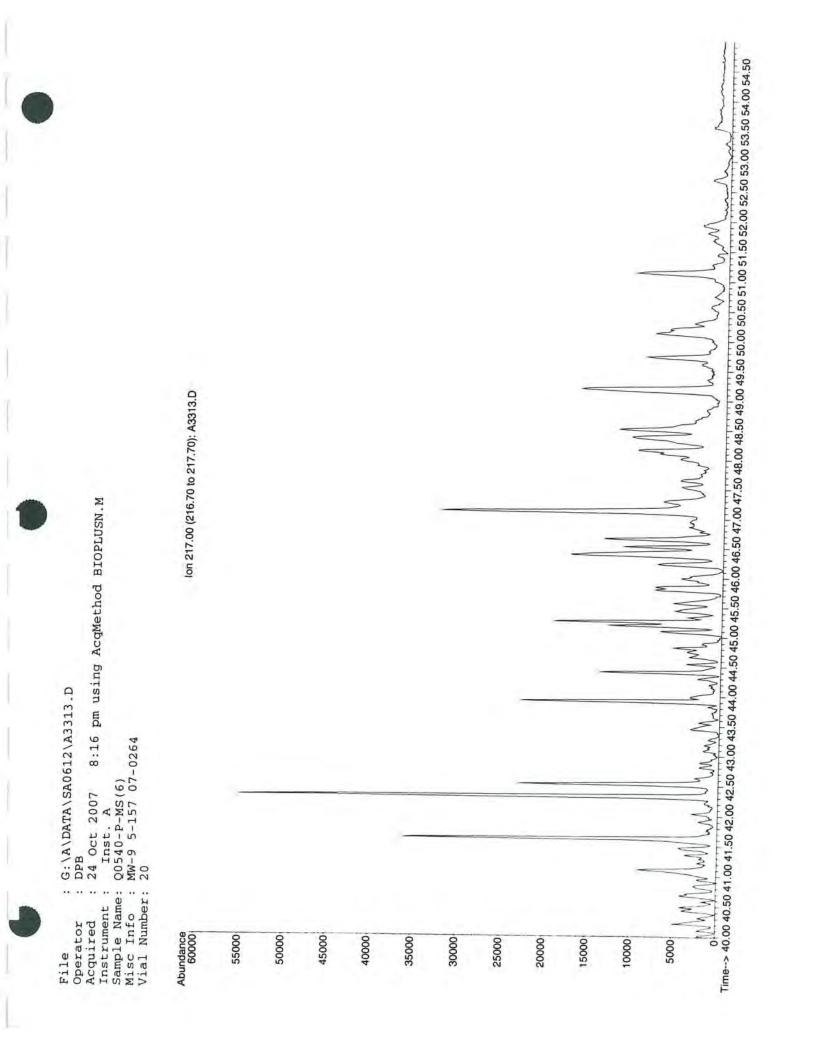
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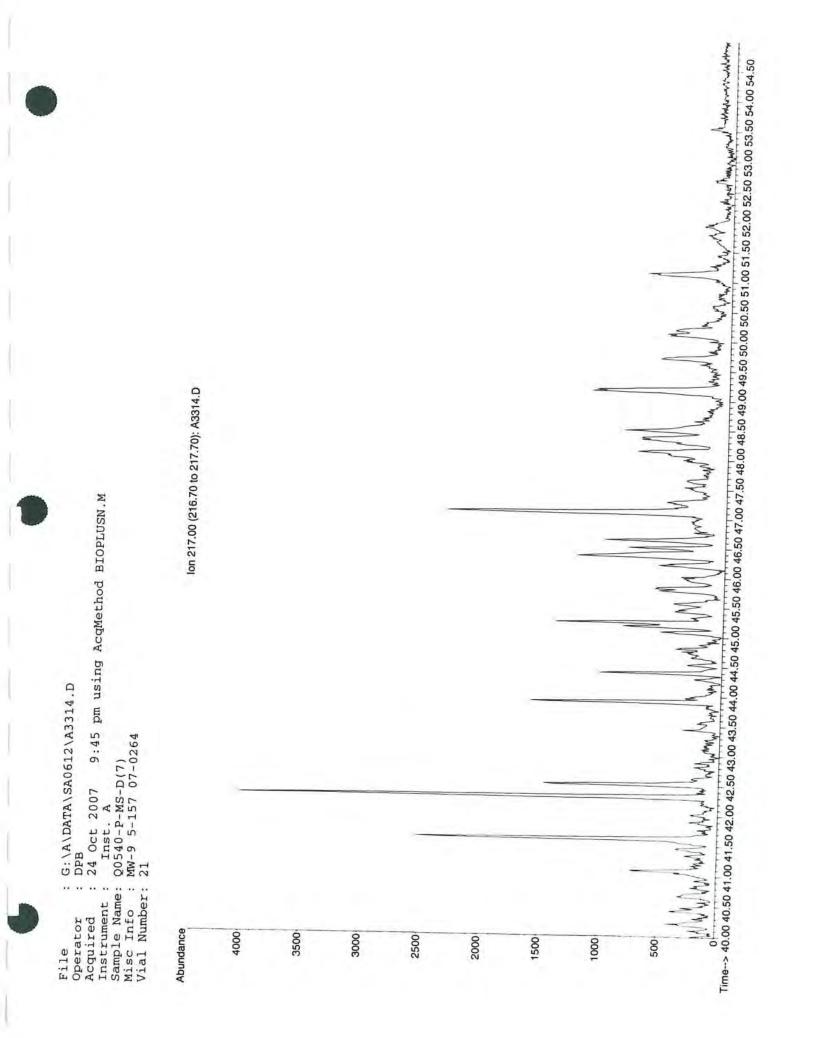
File : G:\A\DATA\SA0612\A3298.I Operator : DPB Acquired : 23 Oct 2007 9:49 pm us Instrument : Inst a		0.8		1121	
		29		EC	
		612\A3		9:49	
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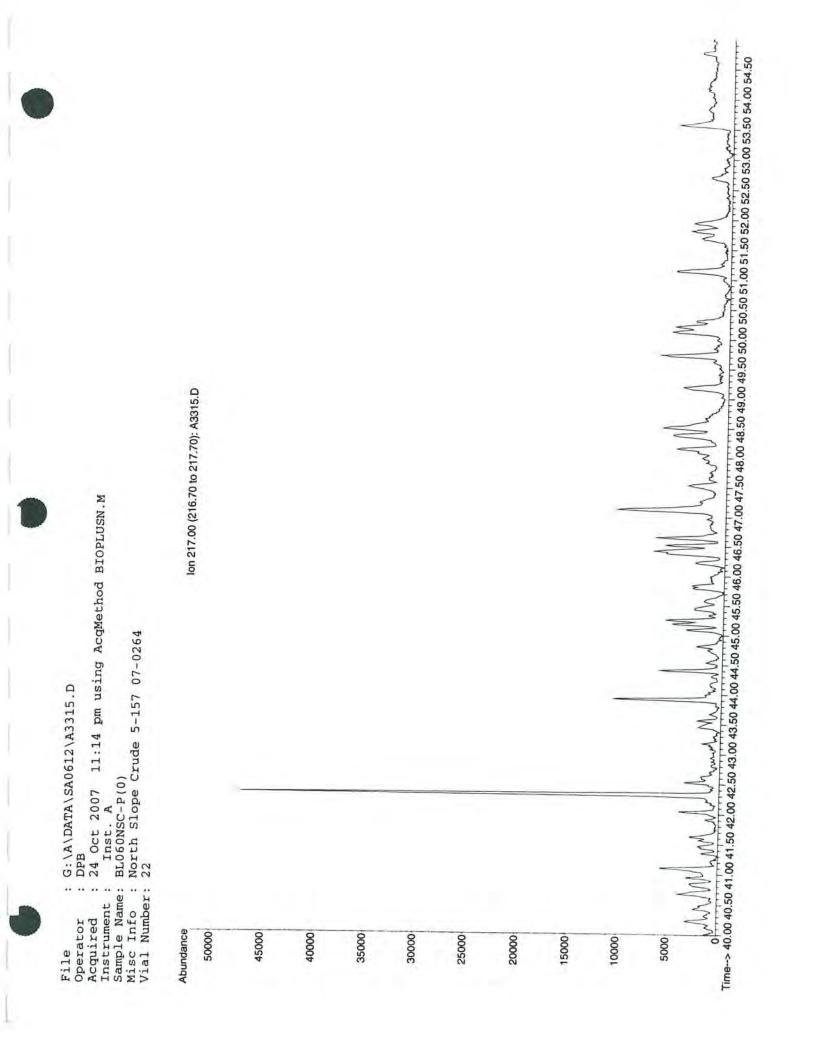
sing AcqMethod BIOPLUSN.M Sample Name: Lnst. A Sample Name: BL058LCS-P-MS(3) Misc Info : Laboratory Control Sample 5-157 07-0264 Vial Number: 5

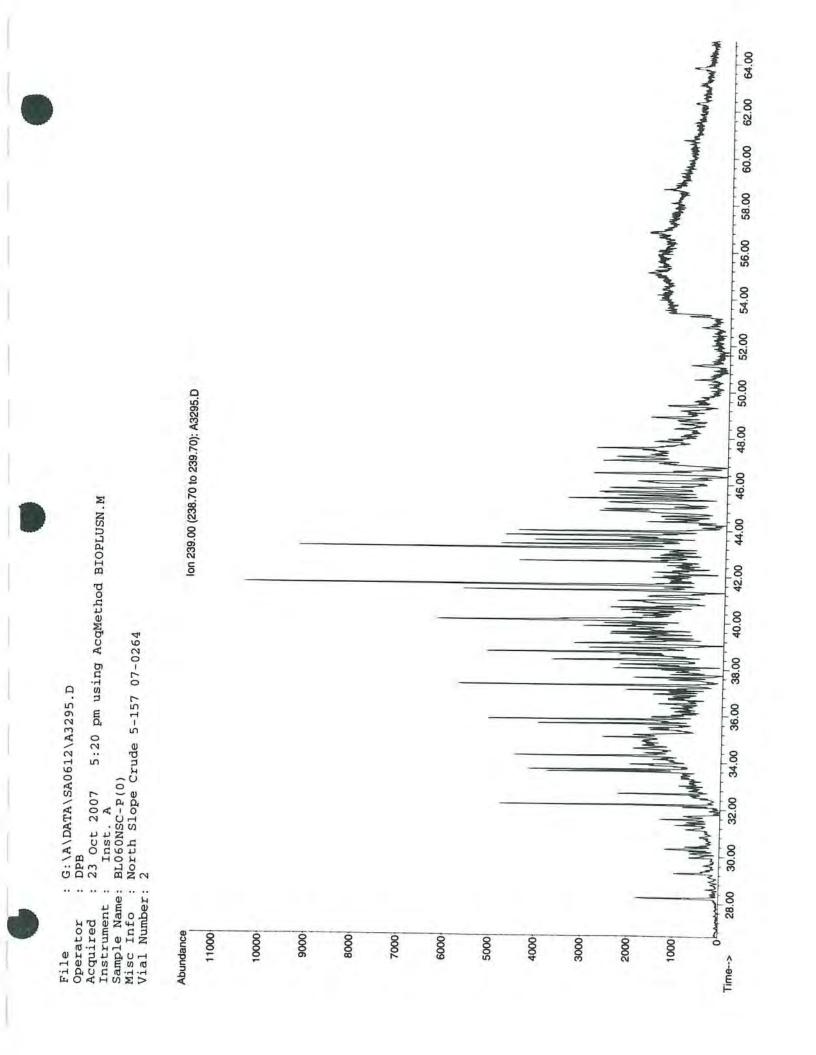
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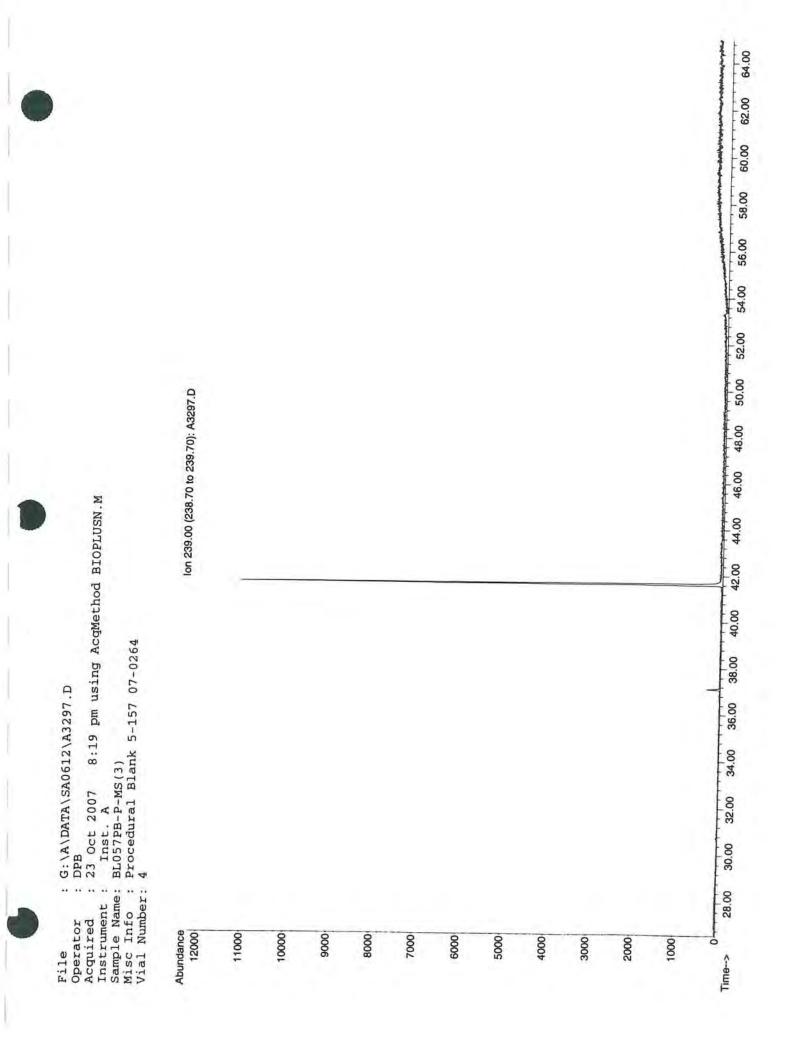


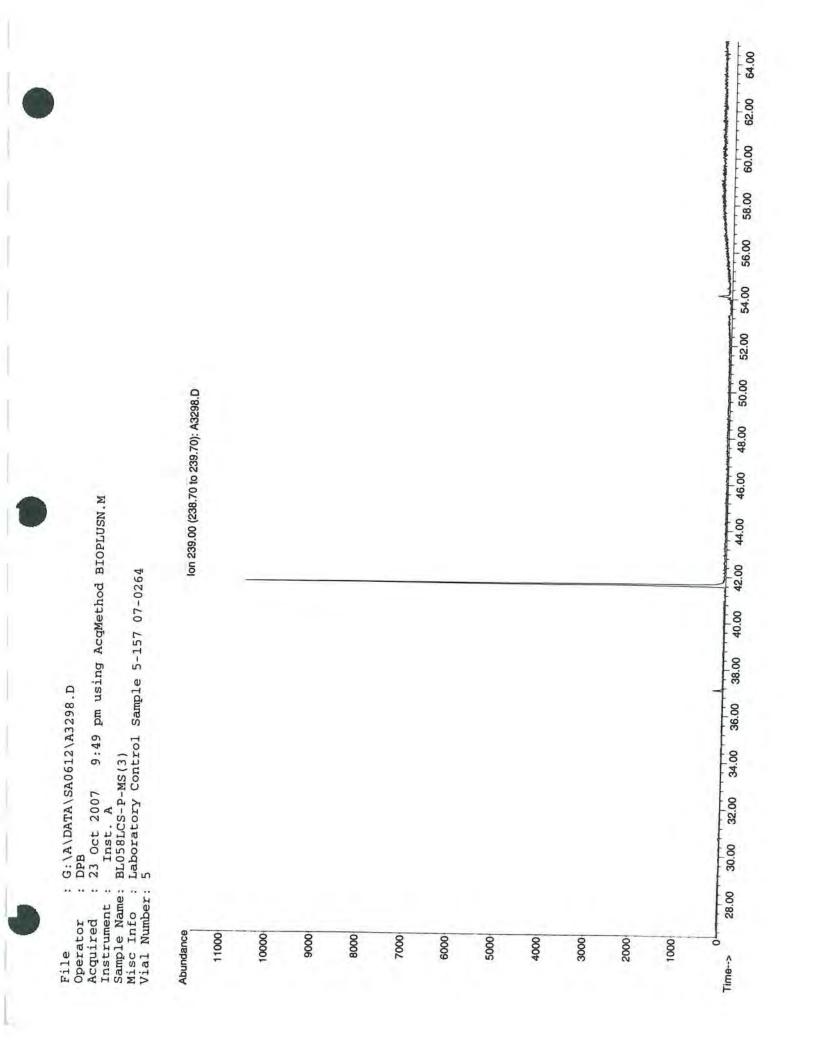


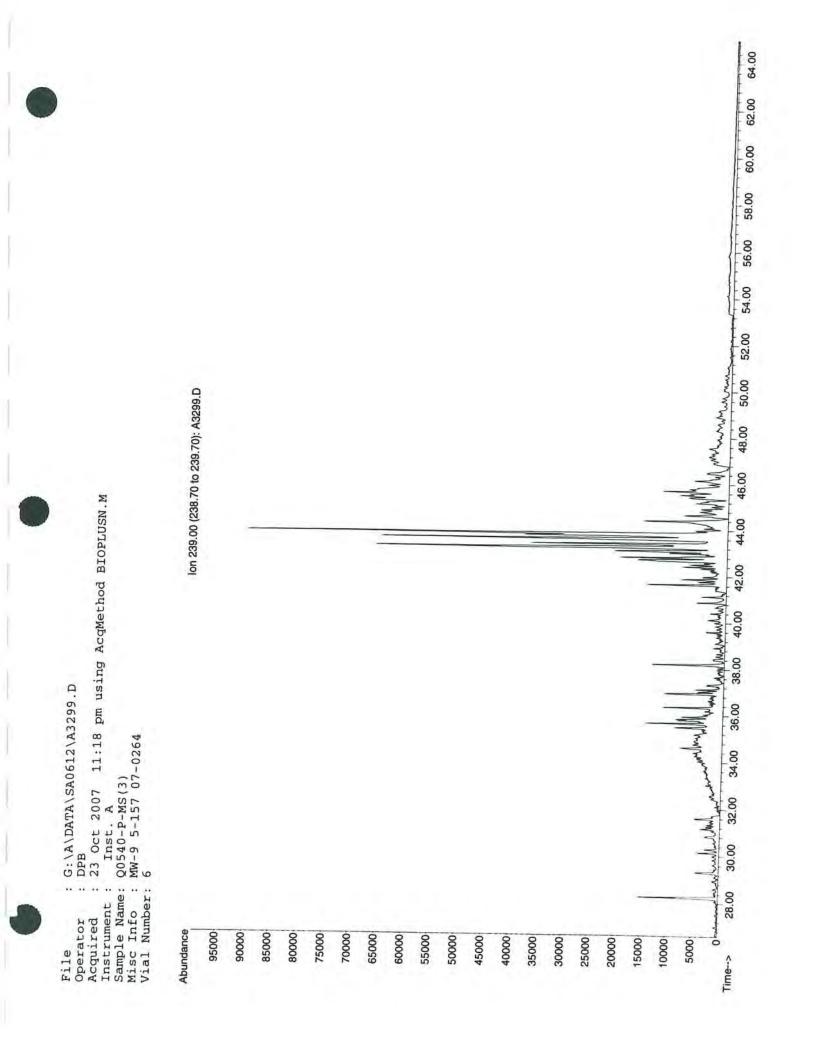


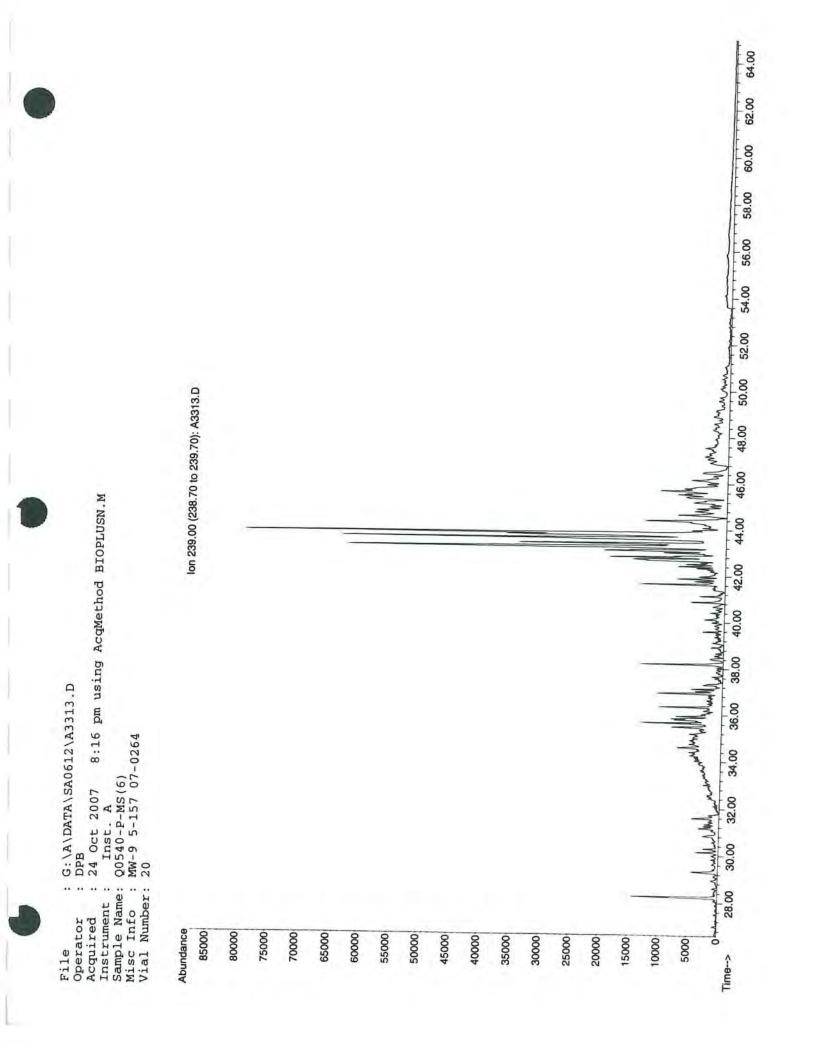


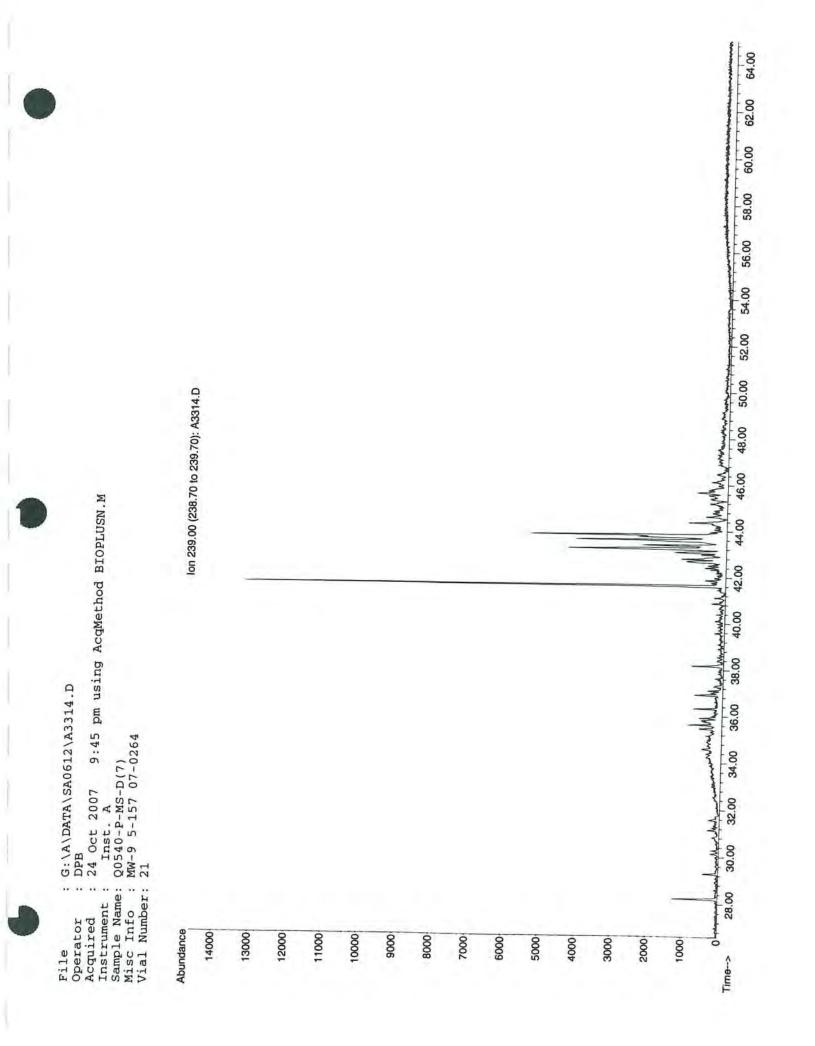


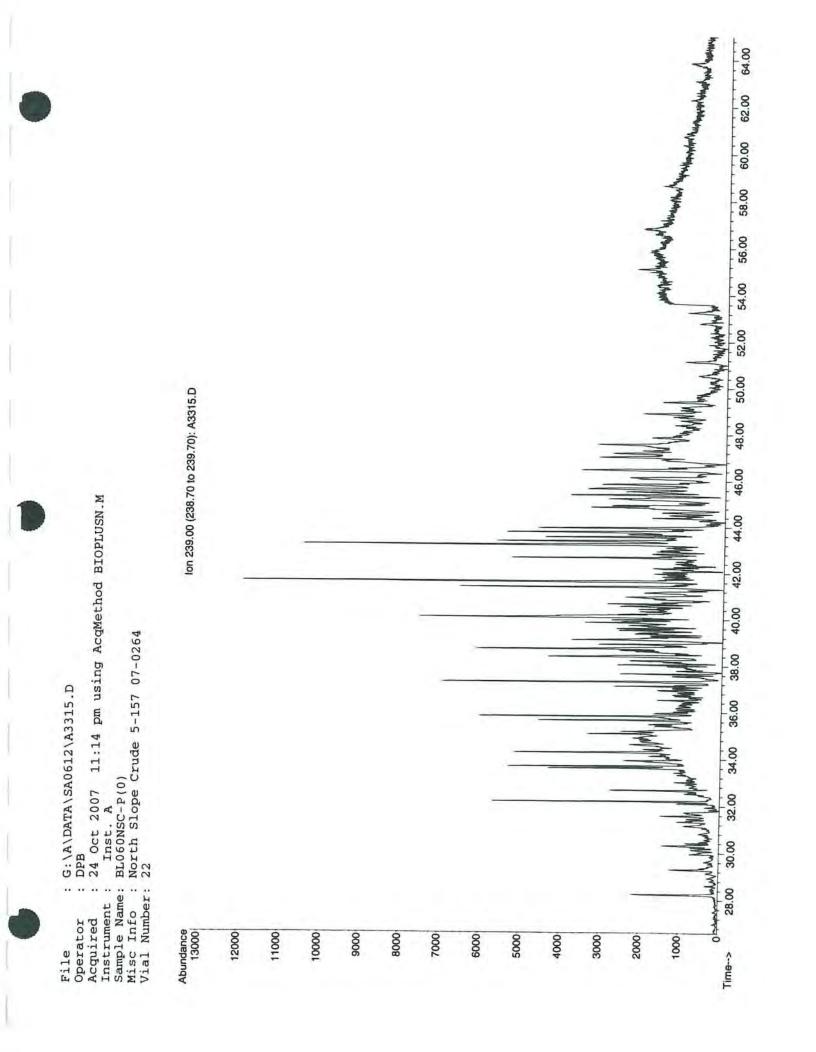


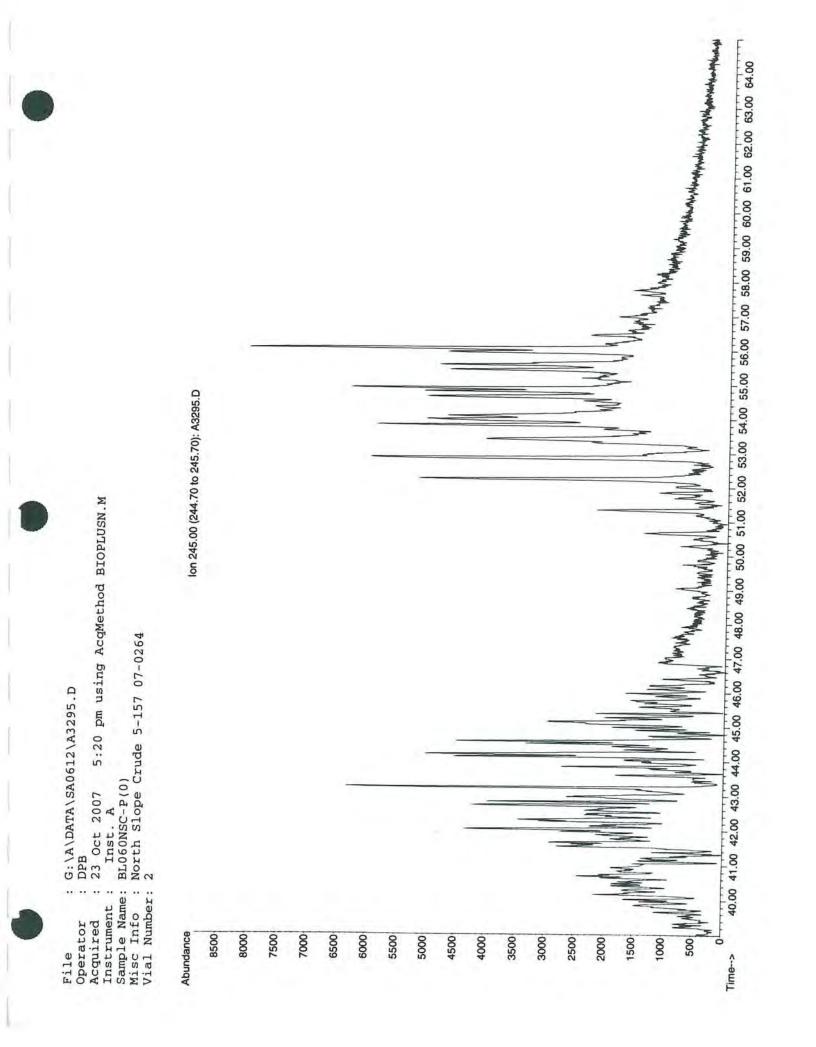


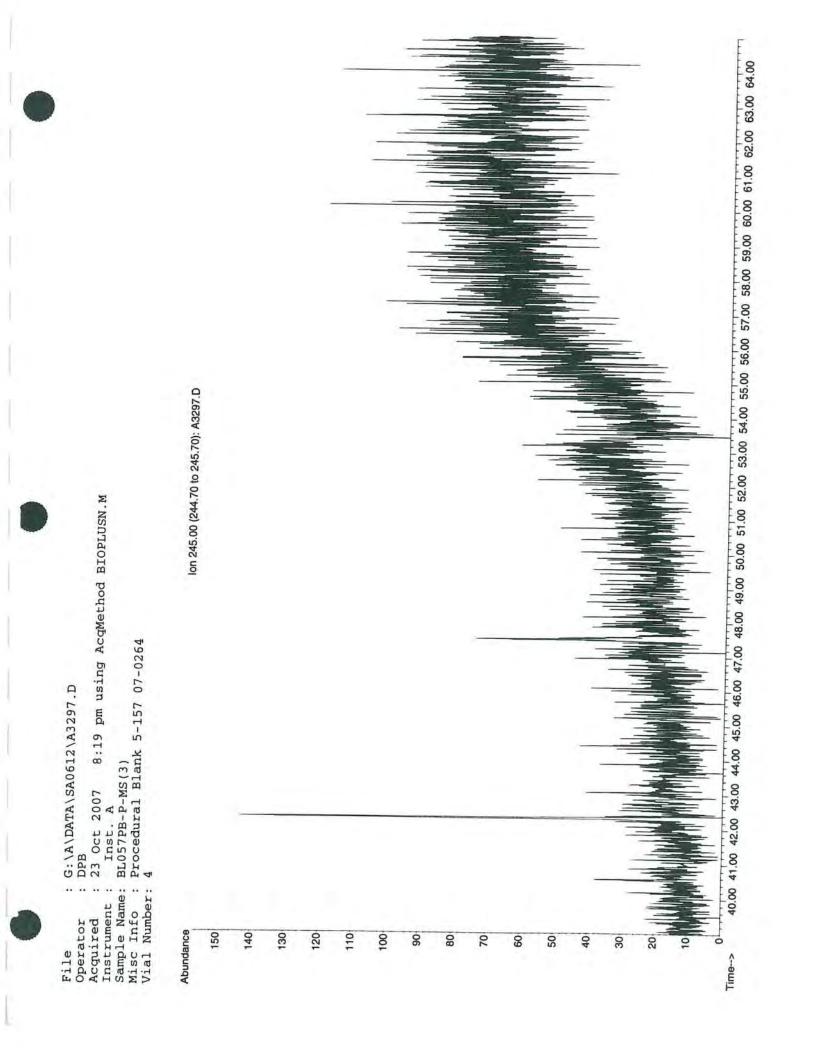


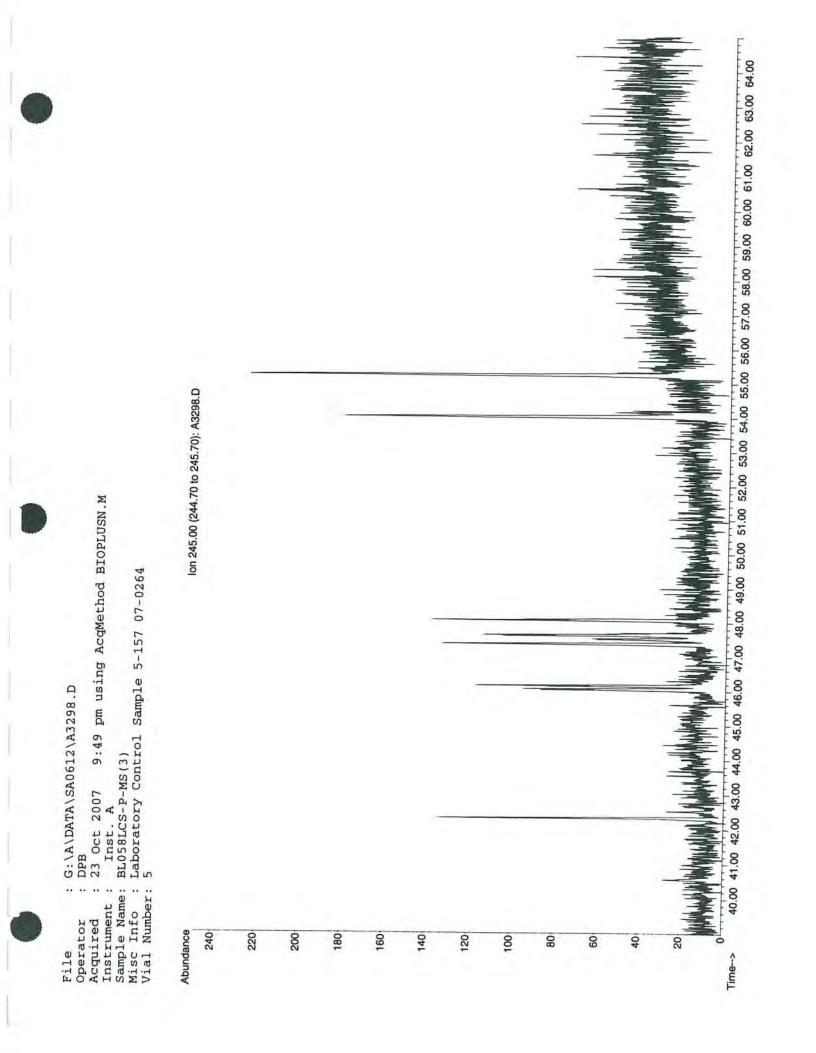


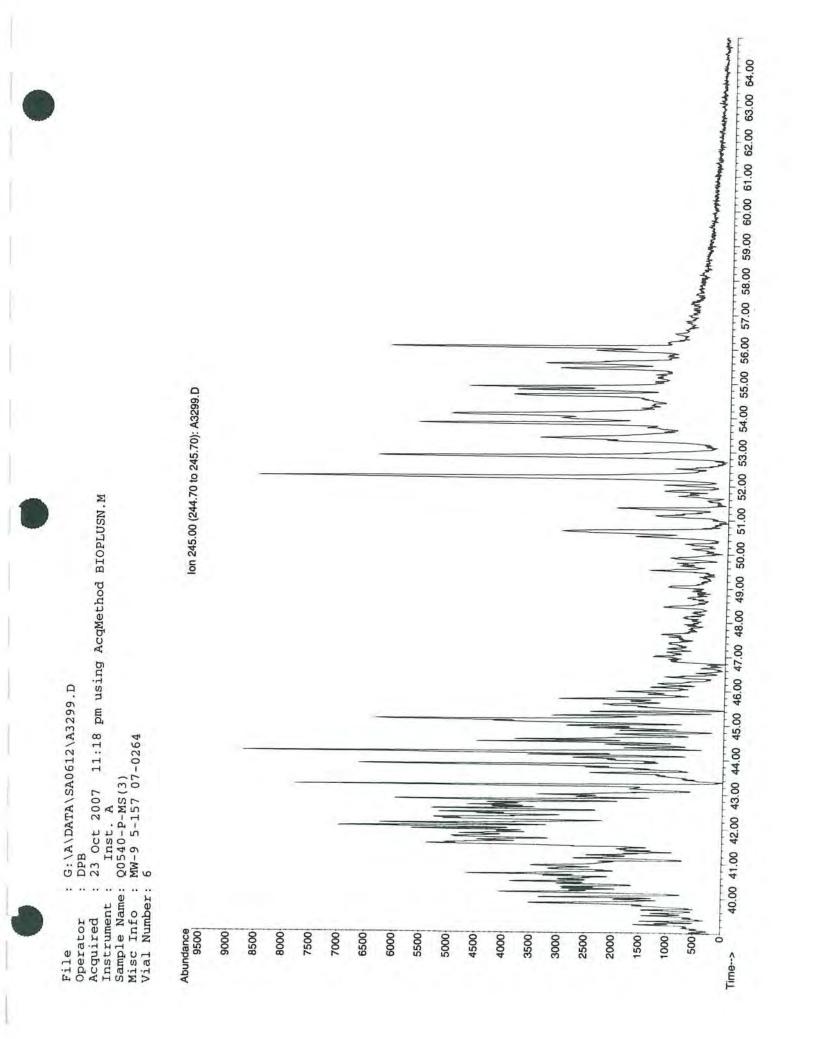


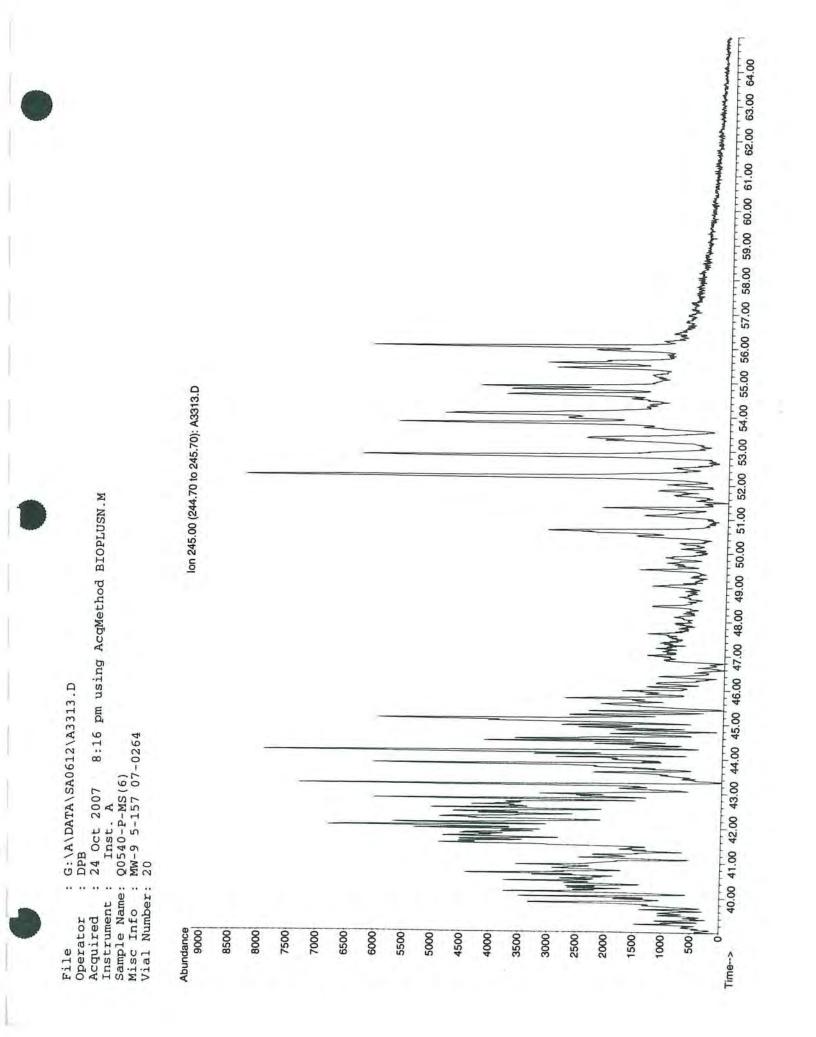


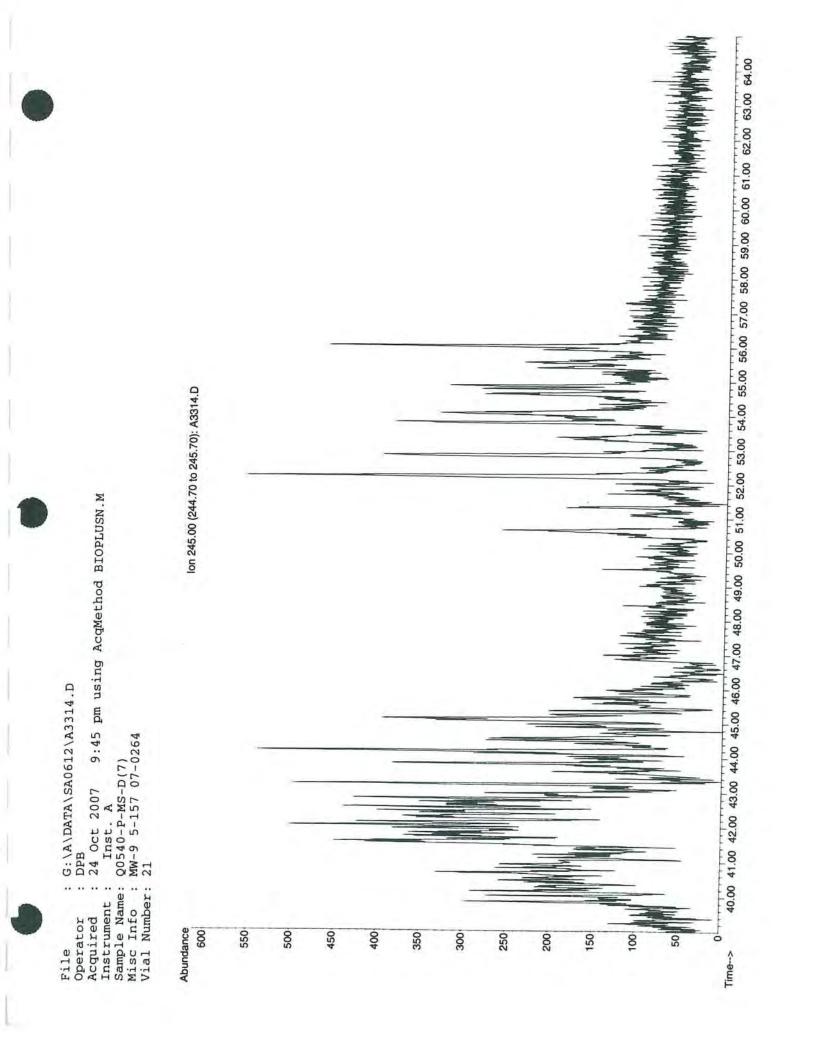


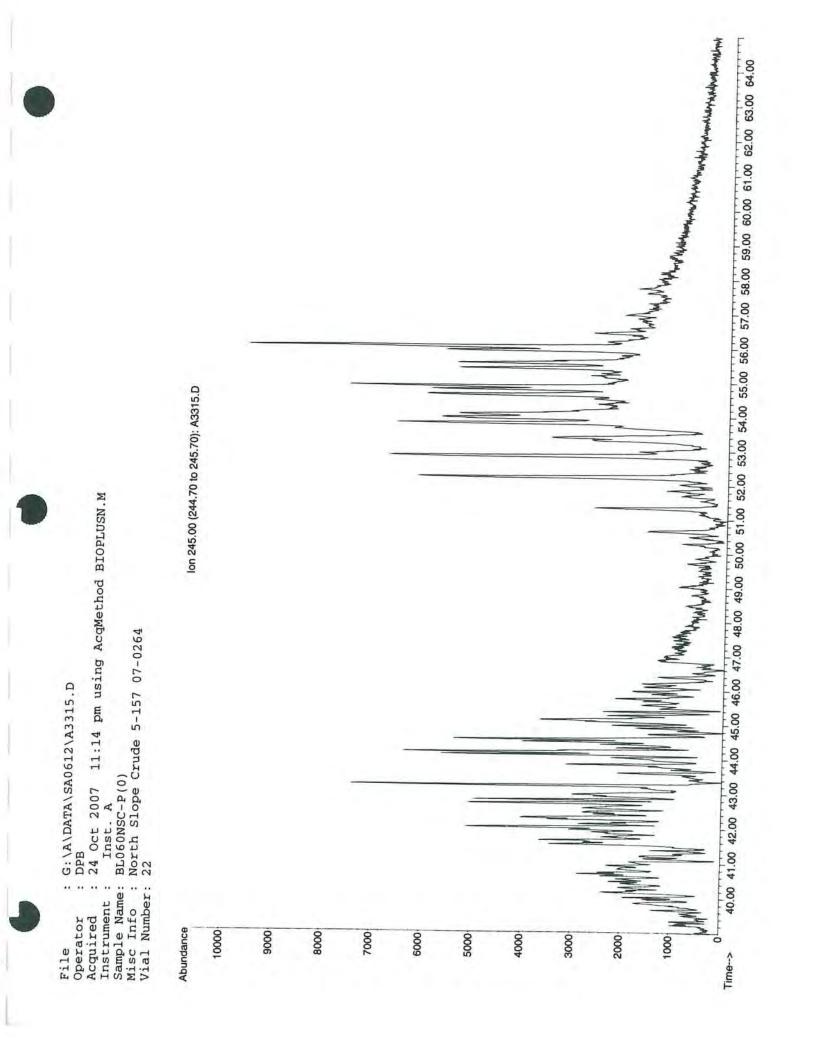












PAH and Biomarker – SEDIMENT QA/QC SUMMARY Batch 07-0266

PROJECT:	Floyd/Snyder – Gas Works Park
PARAMETER:	Polycyclic Aromatic Hydrocarbons (PAH) and Biomarkers
LABORATORY:	Battelle, Duxbury, MA
MATRIX:	Sediment
SAMPLE CUSTODY:	Six soil samples and 1 NAPL were received at Battelle Duxbury Operations (BDO) Laboratory on 10/9/07. Upon Receipt of the samples, the temperature of the cooler was taken and the samples were logged into the laboratory and given unique IDs. The temperature of the cooler upon receipt was slightly below the acceptable range ($4^{\circ}C \pm 2^{\circ}$) at 0.6°C. Also, it was noted that sample Q0540 (MW-9) had some oil residue on the outside of the container as well as the inner side of the bubble wrap. The oil residue was only on the inside of the bubble wrap and did not touch any of the other samples. The client was notified on 10/16/07. The laboratory was instructed to proceed with the analysis. Samples were stored in an access-limited walk-in refrigerator at 4°C until sample preparation could begin.
1	

	Reference Method	Method Blank	Surrogate Recovery	LCS Recovery	MS Recovery	MS/MSD Precision	Control Oil % Diff
SHC and TPH	General NS&T	<5xMDL	40-120% Recovery	40-120% Recovery	40-120% Recovery	≤30% RPD	PD,30% for 90% of the analytes
					MS target spike must be >5x background		

ETHOD:

Soil samples were extracted following general NS&T methods. Approximately 5-8 g of sample was spiked with SHC, PAH, and biomarker surrogates and serial extracted three times with dichloromethane using orbital shaker table techniques. The combined extracts were dried over anhydrous sodium sulfate and concentrated by Kuderna-Danish and nitrogen evaporation techniques. The sample extracts were split in half: one-half of the extract was removed for archiving; the other half was processed through an alumina gravity column to isolate the hydrocarbon fractions of interest. The weights of the resulting extracts were determined gravimetrically. The extracts were concentrated to 1 mL, split, and spiked with internal standard (IS). The pre-injection volume and/or extract split were adjusted to 5mg/mL. One extract was submitted for PAH and the second extract was submitted for SHC and TPH analysis.

PAH and petroleum biomarkers were measured by gas chromatography -mass spectrometry (GC/MS) in the selected ion mode (SIM). An initial calibration consisting of target analytes was completed prior to analysis to demonstrate the linear range of the analysis. Calibration verification was performed at the beginning and end of each 12 hour period in which samples were analyzed. Concentrations of the PAH and petroleum biomarkers were calculated by the internal standard method. Target PAH were quantified using the average RF generated from the initial calibration. The alkyl homologue PAH series were assigned the RF of the parent PAH, Steranes were assigned the RF of cholestane, and triterpanes were assigned the RF of Moretane.

Note: the reporting limit for the alkyl benzene compounds is orders of magnitude higher than the reporting limits for the rest of the PAH compounds.

PAH and Biomarker - SEDIMENT QA/QC SUMMARY Batch 07-0266

PROCEDURAL	
PROCEDURAL BLANK (PB):	A procedural blank (PB) was prepared with each analytical batch. Blanks were analyzed to ensure the sample extraction and analysis methods were free of contamination.
	07-0266 – No exceedences noted.
	Comments – None.
LABORATORY CONTROL SAMPLE (LCS):	A laboratory control sample (LCS) was prepared with each analytical batch. The percent recoveries of target analytes were calculated to measure data quality in terms of accuracy.
	07-0266 – No exceedences noted.
	Comments – None.
SURROGATE	Five surrogate compounds were added prior to extraction, including naphthalene-d8,
RECOVERY:	acenaphthene-d10, phenanthrene-d10, benzo(a)pyrene-d12, and 5b(H)-chloane. The recovery of the surrogate compound was calculated to measure data quality in terms of accuracy (extraction efficiency).
	07-0266 – All four samples extracted in this batch failed SIS recovery criteria with high recoveries.
	Comments – These samples have been extracted twice with similar results. The issue appears to be matrix related rather than extraction related as the PB and LCS (samples without matrix) did not exhibit similar trends.
CONTROL OIL:	A control oil (North Slope Crude) was prepared with the analytical batch. The percent difference (PD) between the measured value and the target value was calculated to measure data quality in terms of accuracy.
	07-0266 – No exceedences noted.
	Comments – None.
CALIBRATIONS:	The GC/MS is collibrated with the state of t
CALIDICATIONS:	The GC/MS is calibrated with a minimum 5 level curve for all compounds. The percent relative standard deviation (% RSD) between RF for the individual target analytes must be $\leq 25\%$, and the mean RSD of all target analytes must be $\leq 15\%$. Each batch of samples analyzed is bracketed by a continuing calibration verification (CCV) sample, run at a frequency of minimally every 12 hours. The PD between the true value and the CCV should be $\leq 25\%$ for individual analytes. Additionally an initial calibration check (ICC) sample is run immediately after each initial calibration. The PD between the ICC and the initial calibration should be $\leq 25\%$

07-0266 - No exceedences noted.

Comments - None.

calibration should be $\leq 25\%$.

Battelle

The Business of Innovation

Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	SB-8 S5 9-10.5	SB-13 2.5-4.0	SB-12A S3 5-6.5	GP-9 7-8
Battelle ID	Q0542-P1	Q0543-P1	Q0544-P1	Q0546-P1
Sample Type	SA	SA	SA	SA
Collection Date	09/18/07	09/20/07	09/20/07	THE REAL PROPERTY AND A DESCRIPTION OF A
Extraction Date	10/23/07	10/23/07	10/23/07	09/18/07
Analysis Date	11/14/07	11/14/07	11/14/07	10/23/07
Analytical Instrument	MS	MS	MS	11/13/07
% Moisture	8.06	16.77		MS
% Lipid	NA	NA	13.98	49.24
Matrix	SOIL		NA	NA
Sample Size	4.78	SOIL	SOIL	SOIL
Size Unit-Basis		4.45	4.63	2.84
Units	G_DRY	G_DRY	G_DRY	G_DRY
Units	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
C3-Alkylbenzenes	3892.89	186.94 J	47632.7	5372.73 J
C4-Alkylbenzenes	18749.55	249.65 J	33828.46	5577.88 J
C5-Alkylbenzenes	5557.54	156.88 J	10399.43 J	
C6-Alkylbenzenes	4402.57	98.64 J	8583.42 J	2089.59 J
Benzo(b)thiophene	3378.11	5076.38		1729.16 J
C1-benzo(b)thiophenes	3329.38		393493.15 D	3568.95
C2-benzo(b)thiophenes	2895.91	1139.53	178498.3 D	3388.9
C3-benzo(b)thiophenes	2534.91	1482.64	145715.34 D	U
		1140.93	63061.83 D	U
24-benzo(b)thiophenes	1632.18	412.32	16930.09 D	U
Naphthalene	197097.18 D	46290.73	4524558.48 D	10239.32
C1-Naphthalenes	98316.58 D	9577.93	1193806.3 D	2301.13
2-Naphthalenes	43038.8 D	8881.07	641883.07 D	6972.28
3-Naphthalenes	20997.44 D	7590.06	260962.9 D	5843.44
24-Naphthalenes	9953 1 D	3506.08	75069.96 D	U
1-Biphenyls + Dibenzofuran	4673.01	7971.01	95177.23	1984.95
liphenyl	9344.63	14005.81	141278.54	856.33
2-Biphenyls + C1-Dibenzolurans	5754.07	4533.09	91796.6	7146.12
cenaphthylene	170.56	23360.5	196999.75	
cenaphthene	5931.38	2216.54	128297.35	79299.94
libenzofuran	1232.99	5571.55		622.88
luorene	2703.28	22655.84	43413.34	454.05
1-Fluorenes	3297.95	5254.37	242645.7	667.43
2-Fluorenes	3895.26		127932.59	U
3-Fluorenes	2966.17	4151.98	86930.23	u
4-Fluorenes		U	41379.34	U
nthracene	2393.37	U	13759.75	U
henanthrene	268.41	88166.17 D	327451.33 D	10159.52
	3271.85	515173.52 D	1652158.57 D	2405.95
1-Phenanthrenes/Anthracenes	5110.85	76738.48 D	713534.16 D	6776.77
2-Phenanthrenes/Anthracenes	5084.27	31628.12 D	351181.43 D	7611.12
3-Phenanthrenes/Anthracenes	2895.42	11745.34 D	103700.95 D	7070.4
4-Phenanthrenes/Anthracenes	1396.86	2593.54 D	23157.55 D	U
etene	2268,57	1860.34	Ŭ	3378.98
ibenzothiophene	561.92	58124.16	248996.12	979.76
1-Dibenzothiophenes	1416.11	15918.52	204065.03	
2-Dibenzothiophenes	1656.13	10229.11	159270.15	2546.1
3-Dibenzothiophenes	1020.41	4900.03		5089.4
4-Dibenzothiophenes	432.74	1443.07	69948.85	4871.82
uoranthene	425.84		19585.16	2457.45
rene	632.53	740795.46 D	769240.56 D	4503.06
1-Fluoranthenes/Pyrenes		954314.48 D	1147697.77 D	14763.15
2-Fluoranthenes/Pyrenes	715.32	120979.11 D	517881 D	13411.77
3-Fluoranthenes/Pyrenes	578	28376.19 D	182979.44 D	19334.13
	463.84	9383.55 D	71821.42 D	U
4-Fluoranthenes/Pyrenes	262.87	U	23035.94	U
)-Benzo(b)naphthothiophenes	45.36	62969.55	105627.53	Ŭ
1-Benzo(b)naphthothiophenes	140.59	19773.48	126444.47	Ŭ
2-Benzo(b)naphthothiophenes	142.96	6070.07	74933.29	ŭ
3-Benzo(b)naphthothiophenes	87.78	7872.43	33192.02	Ŭ
4-Benzo(b)naphthothiophenes	U	725.56	8221.03	U
enzo(a)anthracene	124.37	223728.55 D	290498.83 D	3336.09
hrysene				

Analyzed by Restucci Jr, Richard 11/15/2007

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SB-8 S5 9-10.5	SB-13 2.5-4.0	SB-12A S3 5-6.5	GP-9 7-8
Q0542-P1	Q0543-P1	00544-P1	Q0546-P1
		Contraction of the second s	SA
09/18/07			09/18/07
10/23/07			10/23/07
11/14/07			11/13/07
MS			MS
			49.24
			49.24 NA
			SOIL
			2.84
NG/G_DRY	NG/G_DRY	NG/G_DRY	G_DRY NG/G_DRY
177.77	45327 58	107122.24 0	2502.02
			3503.96
			U
		Very energy	U
			7404.91
			7060.37
			8672.4
			9161.57
			8317.43
			81298.7
			2717.59
			87159.31
	5001002.43	10020314.57	431075.88
136 NME	40	134 NMF	142 NM
142 NME	47		141 NM
145 NME		11 S S 10 S S S S S S S S S S S S S S S	165 NM
135 NME	49	171 NME	132 NM
133 NUME	49		
	Q0542-P1 SA 09/18/07 10/23/07 11/14/07 MS 8.06 NA SOIL 4.78 G_DRY NG/G_DRY 177.77 174.33 136.57 87.49 70.16 76.29 72.83 104.89 31.08 72.35 11.27 79.03 426259.29	Q0542-P1 Q0543-P1 SA SA 09/18/07 09/20/07 10/23/07 10/23/07 11/14/07 11/14/07 MS MS 8.06 16.77 NA NA SOIL SOIL 4.78 4.45 G_DRY G_DRY NG/G_DRY NG/G_DRY NG/G_DRY NG/G_DRY 177.77 45327.58 174.33 13412.49 D 136.57 5542.83 87.49 4208.54 70.16 215988 D 76.29 22285.07 D 72.83 235414.11 D 104.89 290768.98 D 31.08 70186.94 72.35 298934.42 D 11.27 40419.21 79.03 303454.24 D 426259.29 5061002.49 136 NME 40 142 NME 47 145 NME 46	Q0542-P1 Q0543-P1 Q0544-P1 SA SA SA 09/18/07 09/20/07 09/20/07 10/23/07 10/23/07 10/23/07 11/14/07 11/14/07 11/14/07 11/14/07 11/14/07 11/14/07 MS MS MS 8.06 16.77 13.98 NA NA NA SOIL SOIL SOIL 4.78 4.45 4.63 G_DRY G_DRY G_DRY MG/G_DRY NG/G_DRY NG/G_DRY 177.77 45327.58 197122.34 D 174.33 13412.49 D 102039.96 D 136.57 5542.83 42176.85 87.49 4208.54 12791.94 70.16 215988 D 160335.11 76.29 222855.07 D 170225.35 72.83 235414.11 D 195020.96 104.89 290768.98 D 339611.03 D 31.08 70186.94 76194.88

Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	Procedural Blank	
Battelle ID	BL065PB-P	
Sample Type		
Collection Date	PB	
	10/23/07	
Extraction Date	10/23/07	
Analysis Date	11/13/07	
Analytical Instrument	MS	
% Moisture	22.01	
% Lipid	NA	
Matrix	SEDIMENT	
Sample Size	4.20	
Size Unit-Basis	G_DRY	
Units	NG/G_DRY	
C3-Alkylbenzenes	υ	
C4-Alkylbenzenes		
C5-Alkylbenzenes	U	
	U	
C6-Alkylbenzenes	U	
Benzo(b)thiophene	U	
C1-benzo(b)thiophenes	U	
C2-benzo(b)thiophenes	U	
C3-benzo(b)thiophenes	U	
C4-benzo(b)thiophenes	Ű	
Naphthalene	19.41 N	
C1-Naphthalenes	3.75 J	
C2-Naphthalenes	0.75 U	
C3-Naphthalenes		
C4-Naphthalenes	U	
	U	
C1-Biphenyls + Dibenzofuran	U	
Biphenyl	U	
C2-Biphenyls + C1-Dibenzofurans	U	
Acenaphthylene	0.74 J	
Acenaphthene	U	
Dibenzofuran	Ŭ	
Fluorene	U	
C1-Fluorenes	Ŭ	
C2-Fluorenes	Ŭ	
C3-Fluorenes	ŭ	
C4-Fluorenes	Ŭ	
Anthracene		
Phenanthrene	U	
C1-Phenanthrenes/Anthracenes	1.25 J	
22 Phenanthrenes/Anthracenes	U	
22-Phenanthrenes/Anthracenes	U	
C3-Phenanthrenes/Anthracenes	u	
C4-Phenanthrenes/Anthracenes	U	
Retene	u	
Dibenzothiophene	Ŭ	
C1-Dibenzothiophenes	Ū.	
C2-Dibenzothiophenes	ŭ	
C3-Dibenzothiophenes	Ŭ	
C4-Dibenzothiophenes	U U	
luoranthene		
Pyrene	1.01 J	
	1.72 J	
1-Fluoranthenes/Pyrenes	U	
2-Fluoranthenes/Pyrenes	U	
3-Fluoranthenes/Pyrenes	U	
4-Fluoranthenes/Pyrenes	U	
0-Benzo(b)naphthothiophenes	Ŭ	
1-Benzo(b)naphthothiophenes	ũ	
2-Benzo(b)naphthothiophenes	Ŭ	
3-Benzo(b)naphthothiophenes		
4-Benzo(b)naphthothiophenes	U	
enzo(a)anthracene	U	
Chrysene	0.49 J	
	0.75 J	

Client ID	Procedural Blank	
Battelle ID	BL065PB-P	
Sample Type	PB	
Collection Date	10/23/07	
Extraction Date	10/23/07	
Analysis Date	11/13/07	
Analytical Instrument	MS	
% Moisture	22.01	
% Lipid	NA	
Matrix	SEDIMENT	
Sample Size	4.20	
Size Unit-Basis	G_DRY	
Units	NG/G_DRY	
C1-Chrysenes	U	
C2-Chrysenes	U	
C3-Chrysenes	Ŭ	
C4-Chrysenes	Ŭ	
Benzo(b)fluoranthene	ŭ	
Benzo(k)fluoranthene	Ŭ	
Benzo(e)pyrene	U	
Benzo(a)pyrene	ŭ	
Perylene	Ŭ	
Indeno(1,2,3-cd)pyrene	Ŭ	
Dibenz(a,h)anthracene	Ŭ	
Benzo(g,h,i)perylene	1.3 J	
Total PAH	30.42 J	
	612 P.	
Surrogate Recoveries (%)		
Naphthalene-d8	93	
Acenaphthene-d10	87	
Phenanthrene-d10	94	
Benzo(a)pyrene-d12	96	
5b(H)-Cholane	101	

Client ID	060208-03: Sand White Quartz, -50+70					
Battelle ID	DI 0001 00 D	1				
Sample Type	BL066LCS-P					
Collection Date	LCS					
Extraction Date	10/23/07					
Analysis Date	10/23/07					
and the second	11/13/07					
Analytical Instrument % Moisture	MS					
% Lipid	NA					
Matrix	NA					
Sample Size	SEDIMENT					
Size Unit-Basis	NA					
Units	NA NG		Taraat	0/ Deserves	0	
	NO	-	larget	% Recovery	/ Qualifier	
C3-Alkylbenzenes		U				
C4-Alkylbenzenes		Ū				
C5-Alkylbenzenes		Ũ				
C6-Alkylbenzenes		U				
Benzo(b)thiophene	1598.34		2508.50	64		
C1-benzo(b)thiophenes		U	101103			
C2-benzo(b)thiophenes		u				
C3-benzo(b)thiophenes		U				
C4-benzo(b)thiophenes		U				
Naphthalene	1696.05		2500.50	68		
C1-Naphthalenes		U	(A State of the s			
C2-Naphthalenes		U				
C3-Naphthalenes		U				
C4-Naphthalenes		U				
C1-Biphenyls + Dibenzofuran		U				
Biphenyl	1514.31		2504.25	60		
C2-Biphenyls + C1-Dibenzofurans		U				
Acenaphthylene	1796.09		2502,25	72		
Acenaphthene	1813.64		2501.63	72		
Dibenzofuran	1617.51		2504.75	65		
Fluorene	1761.72		2501.38	70		
C1-Fluorenes		U				
C2-Fluorenes C3-Fluorenes		U				
C4-Fluorenes		U				
Anthracene	126.32	U	Summer.			
Phenanthrene	1801.19		2500.38	72		
C1-Phenanthrenes/Anthracenes	1873.41	Sec. 1	2501.25	75		
C2-Phenanthrenes/Anthracenes		U				
C3-Phenanthrenes/Anthracenes		U				
C4-Phenanthrenes/Anthracenes		U				
Retene		UU				
Dibenzothiophene	1649.82	U	2502 50			
C1-Dibenzothiophenes	1049.82	υ	2502.50	66		
C2-Dibenzothiophenes		ŭ				
C3-Dibenzothiophenes		ŭ				
C4-Dibenzothiophenes		U.				
Fluoranthene	1942.61	0	2501.25	78		
Pyrene	1936.25		2500.88	78		
C1-Fluoranthenes/Pyrenes		U	2300.00			
C2-Fluoranthenes/Pyrenes		ŭ				
C3-Fluoranthenes/Pyrenes		ŭ				
C4-Fluoranthenes/Pyrenes		U				
C0-Benzo(b)naphthothiophenes		Ŭ				
C1-Benzo(b)naphthothiophenes		Ŭ				
C2-Benzo(b)naphthothiophenes		ŭ				
C3-Benzo(b)naphthothiophenes		Ŭ				
C4-Benzo(b)naphthothiophenes		ŭ				
Benzo(a)anthracene	1773.04		2500.63	71		
	10.0.2021					

Client ID	060208-03: Sand, White Quartz, -50+70					
Battelle ID	BL066LCS-P					
Sample Type	LCS					
Collection Date	10/23/07					
Extraction Date	10/23/07					
Analysis Date	11/13/07					
Analytical Instrument	MS					
% Moisture	NA					
% Lipid	NA					
Matrix	SEDIMENT					
Sample Size	SEDIMENT					
Size Unit-Basis	NA					
Units	NG		Target 9	& Recovery	Qualifier	
	110	-	Jarger /	a Recovery	Qualmer	
Chrysene	1630.11		2501.00	65		
C1-Chrysenes		U				
C2-Chrysenes		U				
C3-Chrysenes		U				
C4-Chrysenes		U				
Benzo(b)fluoranthene	1727.03		2502.13	69		
Benzo(k)fluoranthene	1847.36		2501.38	74		
Benzo(e)pyrene	1626.79		2503.50	65		
Benzo(a)pyrene	1744.04		2502.00	70		
Perylene	1671.75		2505.13	67		
ndeno(1,2,3-cd)pyrene	1776.5		2501.25	71		
Dibenz(a,h)anthracene	1750.62		2501.38	70		
Benzo(g,h,i)perylene	1808.55		2500.75	72		
Total PAH	36758.39					
Surrogate Recoveries (%)						
Vaphthalene-d8	62					
cenaphthene-d10	61					
henanthrene-d10	66					
Benzo(a)pyrene-d12 ib(H)-Cholane	64					

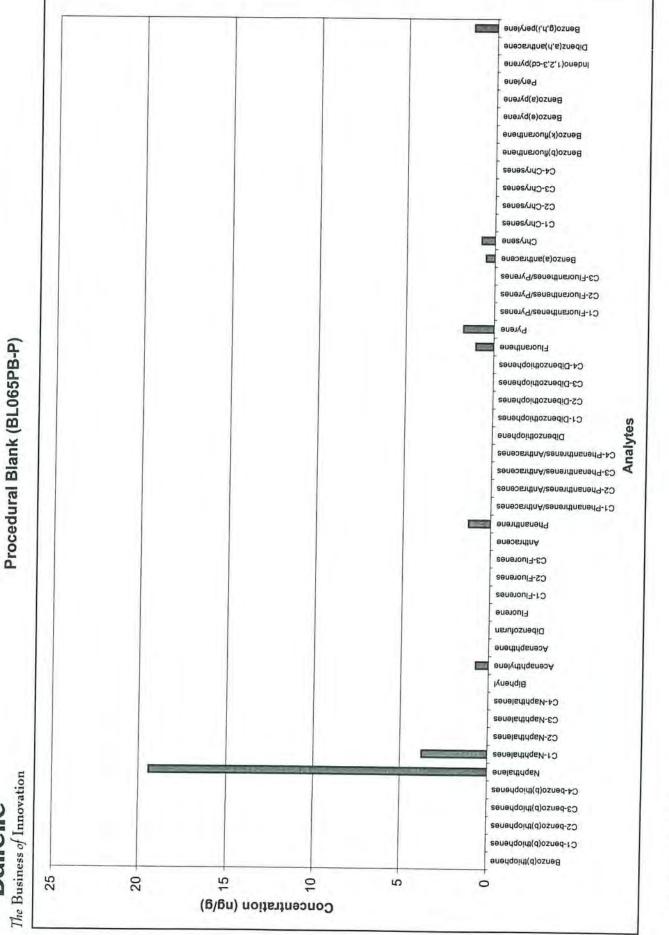
Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	GO98: North Slope Crude				
Battelle ID	DI ASSNICC D				
Sample Type	BL088NSC-P				
Collection Date	NSC				
Extraction Date	10/30/07				
Analysis Date	10/30/07				
Analytical Instrument					
% Moisture	MS				
% Lipid	NA				
Matrix	NA				
Sample Size	OIL				
Size Unit-Basis	5.04				
Units	MG_OIL				
onno.	MG/KG_OIL	Target %	Difference	Qualifier	
C3-Alkylbenzenes	1535.55				
C4-Alkylbenzenes	1217.77				
C5-Alkylbenzenes	694.08				
C6-Alkylbenzenes	443.15				
Benzo(b)thiophene	U				
C1-benzo(b)thiophenes	44.7				
C2-benzo(b)thiophenes	91.01	95.74	4.9		
C3-benzo(b)thiophenes	144.34	132.67			
C4-benzo(b)thiophenes	98.69	96.72	8.8		
Naphthalene	672.77	740.29	2.0		
C1-Naphthalenes	1417.68	1516.04	9.1		
C2-Naphthalenes	1886.29	2000.10	6.5		
C3-Naphthalenes	1497.57		5.7		
C4-Naphthalenes	762.58	1526.96	1.9		
C1-Biphenyls + Dibenzofuran	365.13	898.03	15.1		
Biphenyl	217.33	000.00			
C2-Biphenyls + C1-Dibenzofurans		220.82	1.6		
Acenaphthylene	502.49				
Acenaphthene	8.11	1111			
Dibenzofuran	13.93	14.50	3.9		
Fluorene	68.86	77.75	11.4		
C1-Fluorenes	101.65	92.51	9.9		
C2-Fluorenes	236.93	227.01	4.4		
C3-Fluorenes	324	367.09	11.7		
C4-Fluorenes	235.59	326.32	27.8		
Anthracene	248.33				
Phenanthrene	U	A 14 10			
C1-Phenanthrenes/Anthracenes	272.7	249.49	9.3		
C2-Phenanthrenes/Anthracenes	638.52	549.17	16.3		
C3-Phenanthrenes/Anthracenes	757.4	642,72	17.8		
C4-Phenanthrenes/Anthracenes	537.54	446.11	20.5		
Retene	211.77	180.02	17.6		
Dibenzothiophene	U	1000			
C1-Dibenzothiophenes	228.5	210.35	8.6		
C2-Dibenzothiophenes	466.01	409.03	13.9		
C3-Dibenzothiophenes	651.87	551.46	18.2		
C4-Dibenzothiophenes	526.38	471.36	11.7		
Fluoranthene	315.87	243.11	29.9		
Pyrene	4				
C1-Fluoranthenes/Pyrenes	16.52	12.99	27.2		
C2-Fluoranthenes/Pyrenes	89.15	70.92	25.7		
C3-Fluoranthenes/Pyrenes	150.71	117.89	27.8		
C4-Fluoranthenes/Pyrenes	176.84	137.25	28.8		
CO Banzo/b)coobloatti	133.73				
C0-Benzo(b)naphthothiophenes	56.15				
C1-Benzo(b)naphthothiophenes	215.16				
C2-Benzo(b)naphthothiophenes	277.03				
C3-Benzo(b)naphthothiophenes	179.86				
C4-Benzo(b)naphthothiophenes	125.39				
Benzo(a)anthracene	5.71				

Analyzed by Restucci Jr, Richard 11/15/2007

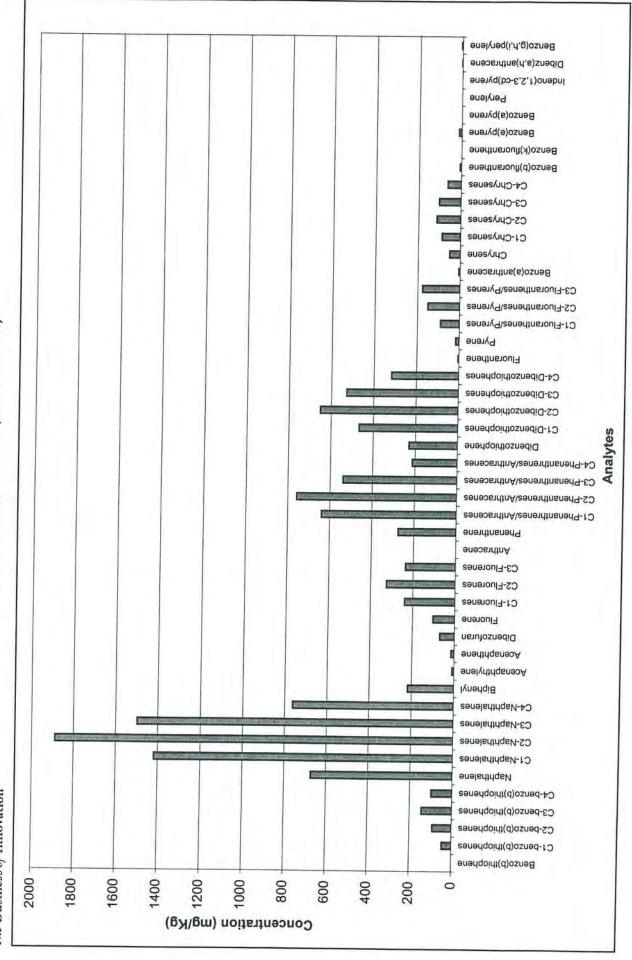
Client ID	GO98: North Slope Crude					
Battelle ID	BL088NSC-P					
Sample Type	NSC					
Collection Date	10/30/07					
Extraction Date	10/30/07					
Analysis Date	11/13/07					
Analytical Instrument	MS					
% Moisture	NA					
% Lipid	NA					
Matrix	OIL					
Sample Size	5.04					
Size Unit-Basis	MG OIL					
Units	MG/KG_OIL	1	Target % D	Difference	Qualifier	
Chrysene	50.54		47.18	7.1		
C1-Chrysenes	86.81		78.82	10.1		
C2-Chrysenes	112.8		102.67	9.9		
C3-Chrysenes	101.95		85.36	19.4		
C4-Chrysenes	62.39		61.99	0.6		
Benzo(b)fluoranthene	5.88		6.08	3.3		
Benzo(k)fluoranthene	0.00	U	0.00	5.5		
Benzo(e)pyrene	12.4	0	12.88	3.7		
Benzo(a)pyrene	12.5	U	12.00	5//		
Perylene		Ű				
ndeno(1,2,3-cd)pyrene		U				
Dibenz(a,h)anthracene	0.8	J				
Benzo(g,h,i)perylene	3.15	1	3.44	8.4		
Total PAH	12929.5		2.2.2	20		
Surrogate Recoveries (%)						
Naphthalene-d8	100					
Acenaphthene-d10	96					
	94					
benanthrong d10						
Phenanthrene-d10 Benzo(a)pyrene-d12	97 105					



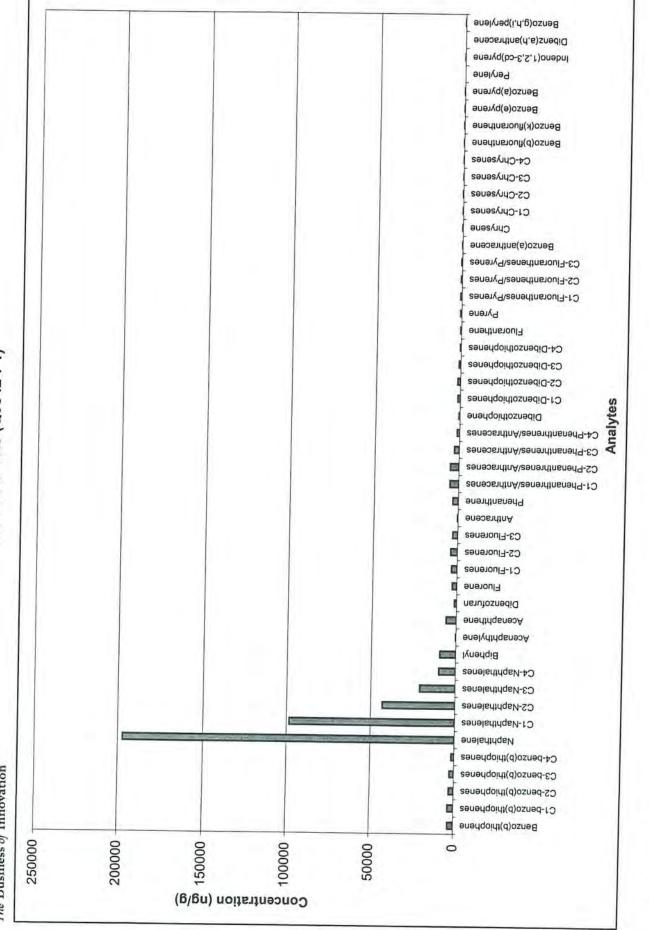


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GO98: North Slope Crude (BL088NSC-P)

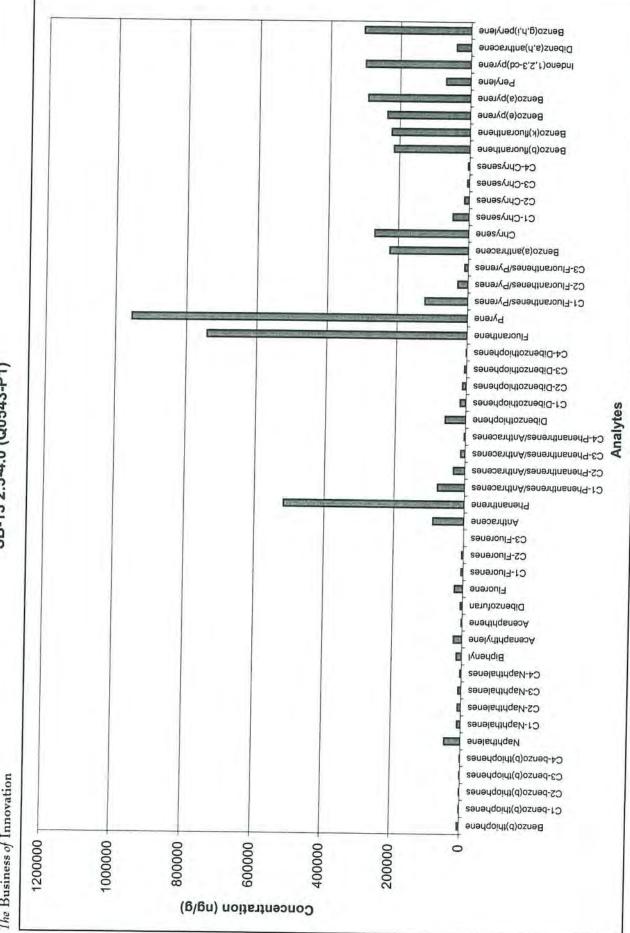


SB-8 S5 9-10.5 (Q0542-P1)



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SB-13 2.5-4.0 (Q0543-P1)



Battelle

SB-12A S3 5-6.5 (Q0544-P1)

					Dipensofusion Private Procession Fluorenci Pluorenci C2-Fluorenci C3-Fluorenci C3-Fluorenci C3-Fluorenci C3-Fluorenthrenes/Anthrescenci C3-Fluorenthrenes/Anthrescenci C4-Phenanthrenacytricenci C4-Phenanthrenacytricenci C4-Phenanthrenacytricenci C3-Fluorenthrenacytricenci C4-Phenanthrenacytricenci C4-Phenanthrenacytricenci C4-Phenanthrenacytricenci C4-Phenanthrenacytricenci C4-Phenanthrenacytricenci C4-Phenanthrenacytricenci C4-Phenanthrenacytricenci C4-Phenanthrenacytricenci C4-Phenanthrenacytricenci C4-Physenes C4-Chr
4500000	4000000	2500000 ng 3000000 ng 2500000 ng 25000000 ng 2500000000 ng 250000000 ng 25000000 ng 250000000 ng 25000000 ng 250000000 ng 25000000000 ng 25000000000 ng 2500000000 ng 250000000000 ng 25000000000000 ng 250000000000000 ng 25000000000000000000000000000000000000	1500000	500000	Benzo(b)(hiophene C1-benzo(b)(hiophene C2-benzo(b)(hiophene C3-benzo(b)(hiophene C3-benzo(b)(hiophene C3-benzo(b)(hiophene C3-hephthalene C3-haphthalene C3-haphthalene C3-haphthalene C3-haphthalene Biphen) Biphen)

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GP-9 7-8 (Q0546-P1)

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C3-Elinorenes	C3-Fluorenes Anthracenes Phenanthrenes/Anthracenes C1-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes C3-Phenanthrenes/Anthracenes	Anthracene Phenanthrenes/Anthracenes Phenanthrenes/Anthracenes D4-Phenanthrenes/Anthracenes Dibenzothiophenes C2-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes C3-Dibenzothiophenes							20010					C4-benzo(b)thiophenes Waphthalenes C1-Waphthalenes C3-Waphthalenes C4-Waphthalenes Biphenyl Acenaphthylene Pibenzofuran C1-Fluorenes C1-Fluorenes C3-Fluorenes
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Project Client: Floyd|Snider Project Name: Gas Works Park Project Number: N007097-0001

Client ID	SB-8 S5 9-10.5	SB-13 2.5-4.0	SB-12A S3 5-6.5	GP-97-8
Battelle ID	Q0542-P1	Q0543-P1	00544 54	a survey and
Sample Type	SA	Q0543-P1 SA	Q0544-P1	Q0546-P1
Collection Date	09/18/07		SA	SA
Extraction Date	10/23/07	09/20/07	09/20/07	09/18/07
Analysis Date	11/14/07	10/23/07	10/23/07	10/23/07
Analytical Instrument	MS	11/14/07	11/14/07	11/13/07
% Moisture	8.06	MS	MS	MS
% Lipid	NA	16.77	13.98	49.24
Matrix	SOIL	NA	NA	NA
Sample Size		SOIL	SOIL	SOIL
Size Unit-Basis	4.78	4.45	4.63	2.84
Units	G_DRY	G_DRY	G_DRY	G_DRY
Units .	NG/G_DRY	NG/G_DRY	NG/G_DRY	NG/G_DRY
C23 Tricyclic Terpane	162.73	U	809.73	
C29 Tricyclic Terpane -22S	27.09	Ŭ		U
C29 Tricyclic Terpane -22R	25	ŭ	314.6	596.59
18a(H)-22,29,30-Trisnorneohopane -TS	17.34 J	U	326.97	722.25
17a(H)-22,29,30-Trisnorhopane -TM	32.61	34.85 J	229.08	U
30-Norhopane	94.28		483.22	U
18a(H) & 18b(H)-Oleananes	35.41	73.47 J	1399.33	1551.53
Hopane		70.63 J	333.67	596.7
30-Homohopane -22S	164.54 66.91	151.58	1994.84	1270.93
30-Homohopane -22R	37.11	61.44 J	1003.25	896.47
13b(H),17a(H)-20S-Diacholestane		U	630.21	716.91
13b(H),17a(H)-20R-Diacholestane	153.15	187.6	880.4	1117.57
14a(H),17a(H)-20R-methylcholestane	85.75	76.89 J	496.59	3573.01
	209.35	71.97 J	1752.21	2459.83
14a(H),17a(H)-20S-Ethylcholestane 14a(H),17a(H)-20R-Ethylcholestane	133.71	U	U	U
C21-TAS	134.38	70.72 J	1260.44	1428.5
C26-TAS(20S)	37.97	398.7	308.64	590.98
C26.C27-TAS	43.29	U	968.53	343.78
C27-TAS(20R)	129.61	28.11 J	3548.47	3304.51
C28-TAS(20S)	64.93	8.84 J	2175.84	794.95
C28-TAS(20R)	36.12	8.61 J	906.06	1410.9
C21-MAS	32.67	9.93 J	845.84	368.27
C22-MAS	50.35	5.24 J	82.76 J	854.6
C27-MAS	20.34	U	54.36 J	820.26
C27-20R-MAS	15.71	86.56	936.12	U
C27-20S-MAS	41.57	1949.58	3176.64	U
C28-20S-MAS	35.96	440.19	1190.74	276.54
C27-C2920S/R-MAS	73.31	91.98	2517.57	2212.9
C29-20S-MAS	61.6	9478.41	10891.68	1345.44
C29-20S-MAS C29-20R-MAS	27.19	11591.31	U	3971.75
	61.79	1060.24	1105.64	1000.84
TAS_245	U	U	U	U
MAS_239	U	U	U	U
Surrogate Recoverles (%)				
Naphthalene-d8	136 NME			
Acenaphthene-d10	142 NME	40	134 NME	142 NME
Phenanthrene-d10		47	260 NME	141 NME
Benzo(a)pyrene-d12	145 NME	46	152 NME	165 NME
5b(H)-Cholane	135 NME	49	171 NME	132 NME
and the subliding.	155 NME	712 NME	6100 NME	255 NME

Analyzed by Restucci Jr, Richard 11/15/2007

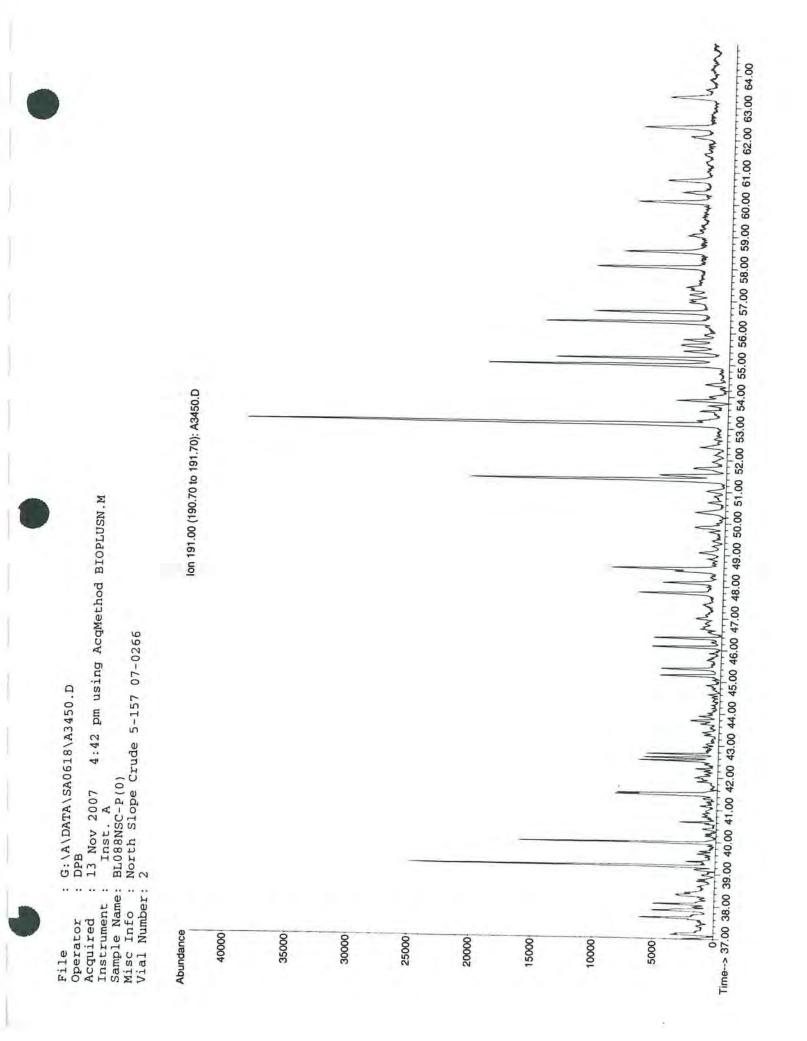
Client ID	Procedural Blank	
Battelle ID	BL065PB-P	
Sample Type		
Collection Date	PB	
	10/23/07	
Extraction Date	10/23/07	
Analysis Date	11/13/07	
Analytical Instrument	MS	
% Moisture	22.01	
% Lipid	NA	
Matrix		
	SEDIMENT	
Sample Size	4.20	
Size Unit-Basis	G_DRY	
Units	NG/G_DRY	
C23 Tricyclic Terpane	U	
C29 Tricyclic Terpane -22S	U	
C29 Tricyclic Terpane -22R	Ŭ	
18a(H)-22,29,30-Trisnomeohopane -TS		
17a/HI-22 20 20 Trisportionage The	U	
17a(H)-22,29,30-Trisnorhopane -TM	U	
30-Norhopane	U	
18a(H) & 18b(H)-Oleananes	U	
Hopane	U	
30-Homohopane -22S	U	
30-Homohopane -22R	U	
13b(H).17a(H)-20S-Diacholestane	Ū	
13b(H), 17a(H)-20R-Diacholestane	ũ	
14a(H),17a(H)-20R-methylcholestane	ŭ	
14a(H),17a(H)-20S-Ethylcholestane	Ŭ	
14a(H),17a(H)-20R-Ethylcholestane		
	U	
C21-TAS	U	
C26-TAS(20S)	U	
C26,C27-TAS	U	
C27-TAS(20R)	U	
C28-TAS(20S)	Ú	
C28-TAS(20R)	Ū	
C21-MAS	Ŭ	
C22-MAS	ŭ	
C27-MAS	U	
C27-20R-MAS		
그렇고 집에 가장 있다. 신가지 않아 봐야?	U	
C27-20S-MAS	Û	
C28-20S-MAS	U	
C27-C2920S/R-MAS	U	
C29-20S-MAS	U	
C29-20R-MAS	U	
TAS_245	U	
MAS_239	Ŭ	
Surrogate Recoveries (%)		
Naphthalene-d8	93	
Acenaphthene-d10	87	
Phenanthrene-d10		
Benzo(a)pyrene-d12	94	
5b(H)-Cholane	96	
su(n)-cholane	101	

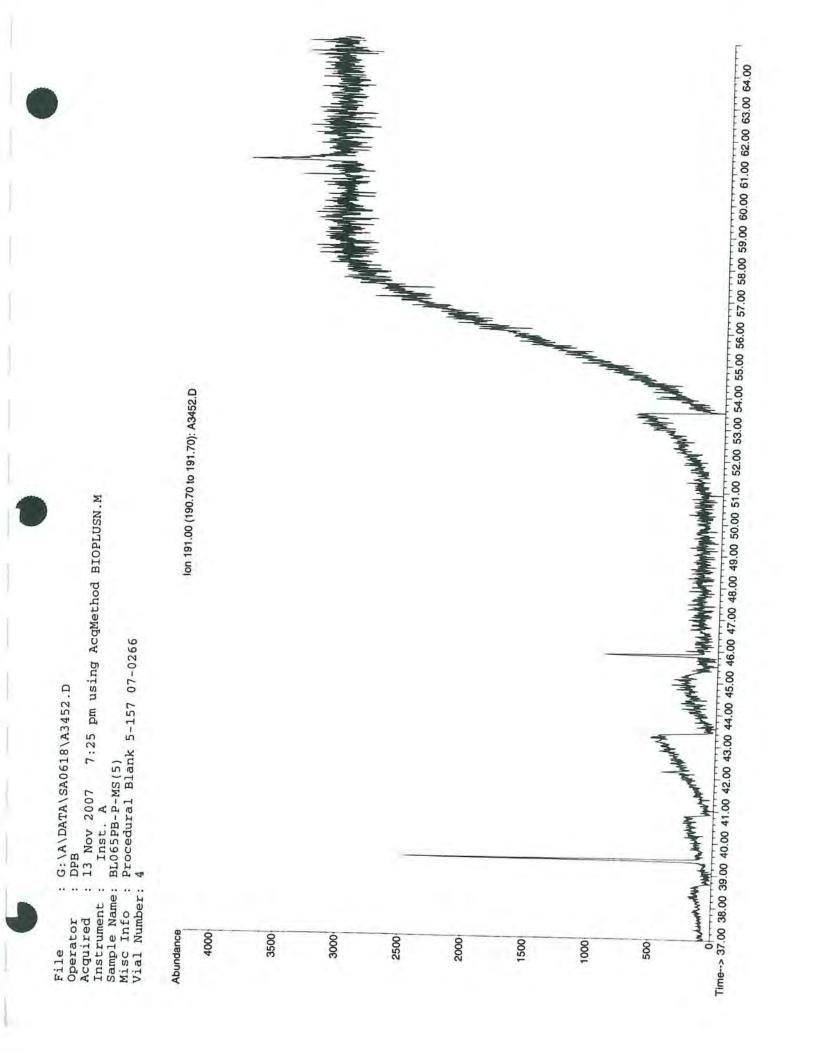
Client ID	060208-03: Sand, White Quartz, -50+70			
Battelle ID	D) 0001 00 0			
Sample Type	BL066LCS-P			
	LCS			
Collection Date	10/23/07			
Extraction Date	10/23/07			
Analysis Date	11/13/07			
Analytical Instrument	MS			
% Moisture	NA			
% Lipid	NA			
Matrix	SEDIMENT			
Sample Size	NA			
Size Unit-Basis	NA			
Units	NG	Target % Recovery	Qualifier	
C23 Tricyclic Terpane				
	U			
C29 Tricyclic Terpane -22S	U			
C29 Tricyclic Terpane -22R	U			
18a(H)-22,29,30-Trisnomeohopane -TS	υ			
17a(H)-22,29,30-Trisnorhopane -TM	U			
30-Norhopane	U			
18a(H) & 18b(H)-Oleananes	U			
Hopane	U			
30-Homohopane -22S	U			
30-Homohopane -22R	U			
13b(H),17a(H)-20S-Diacholestane	U			
13b(H),17a(H)-20R-Diacholestane	U			
14a(H),17a(H)-20R-methylcholestane	U			
14a(H),17a(H)-20S-Ethylcholestane	Ū			
14a(H),17a(H)-20R-Ethylcholestane	Ŭ			
C21-TAS	Ŭ			
C26-TAS(20S)	Ū			
C26,C27-TAS	Ũ			
C27-TAS(20R)	Ŭ			
C28-TAS(20S)	Ŭ			
C28-TAS(20R)	Ŭ			
C21-MAS	Ŭ			
C22-MAS	Ū			
C27-MAS	Ũ			
C27-20R-MAS	Ŭ			
C27-20S-MAS	Ŭ			
C28-205-MAS	ũ			
C27-C2920S/R-MAS	ŭ			
C29-20S-MAS	ŭ			
C29-20R-MAS	ŭ			
TAS_245	Ŭ			
MAS_239	Ŭ			
Surrogate Recoveries (%)				
Naphthalene-d8	62			
Acenaphthene-d10	61			
Phenanthrene-d10				
Benzo(a)pyrene-d12	66			
	64 62			
5b(H)-Cholane				

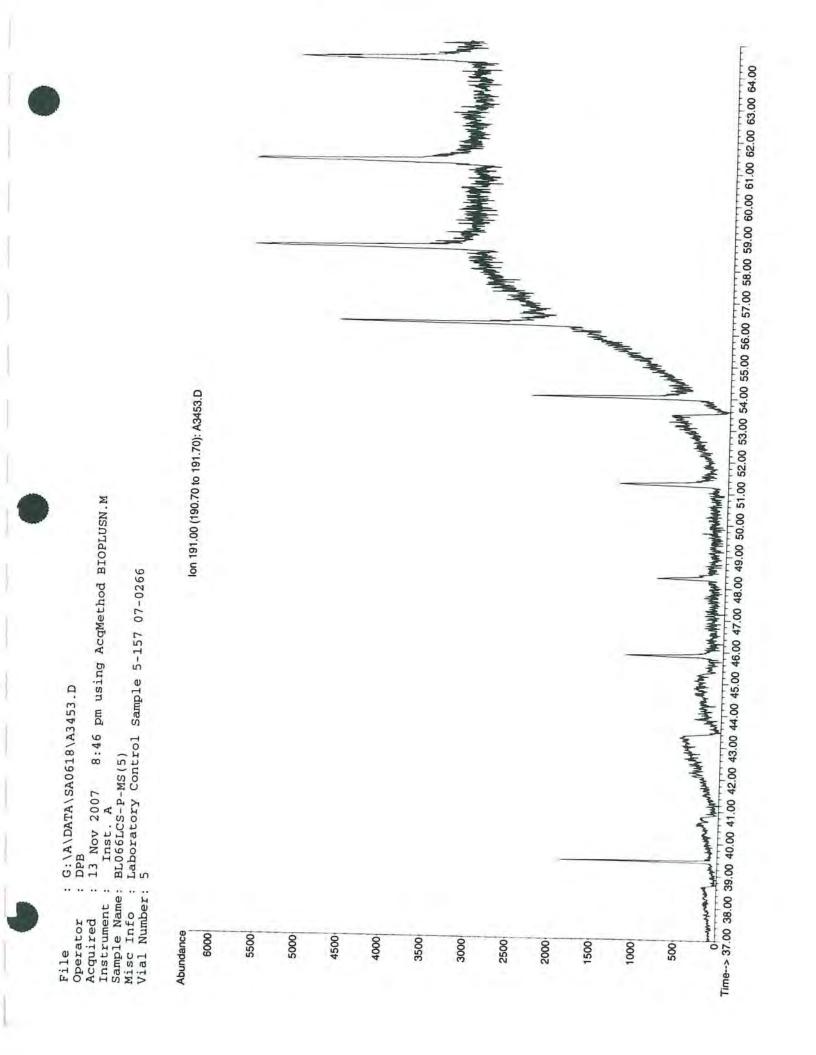
Battelle

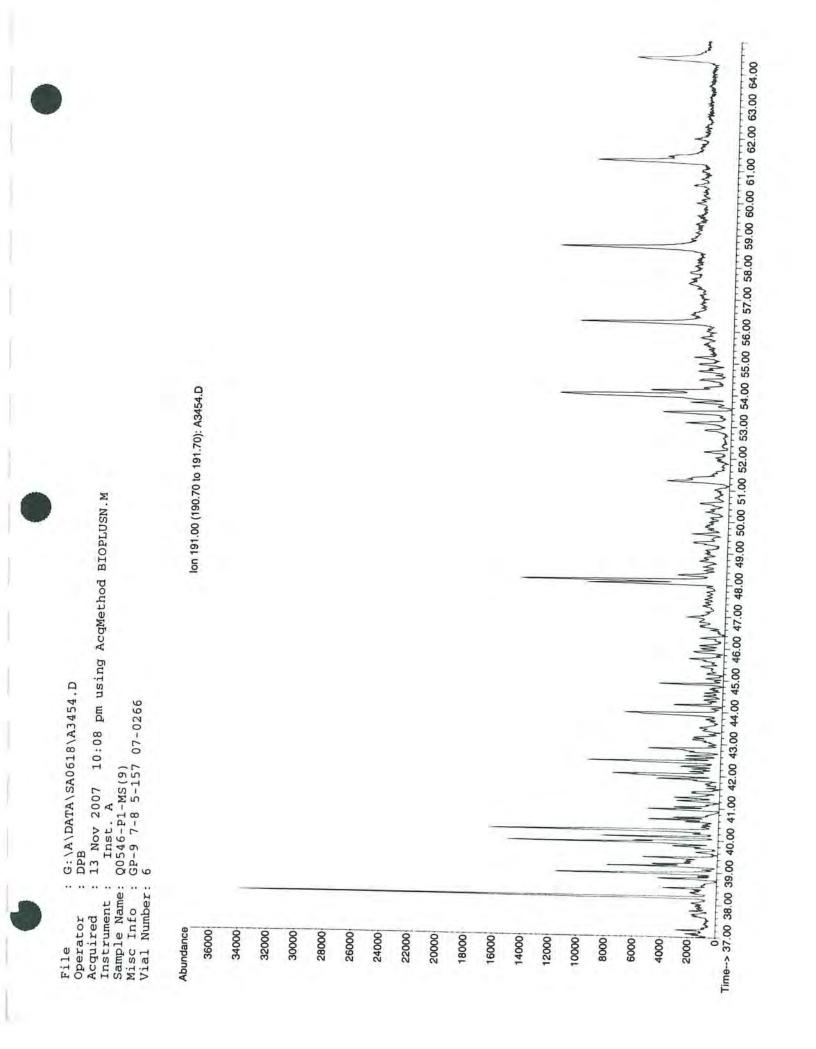
The Business of Innovation

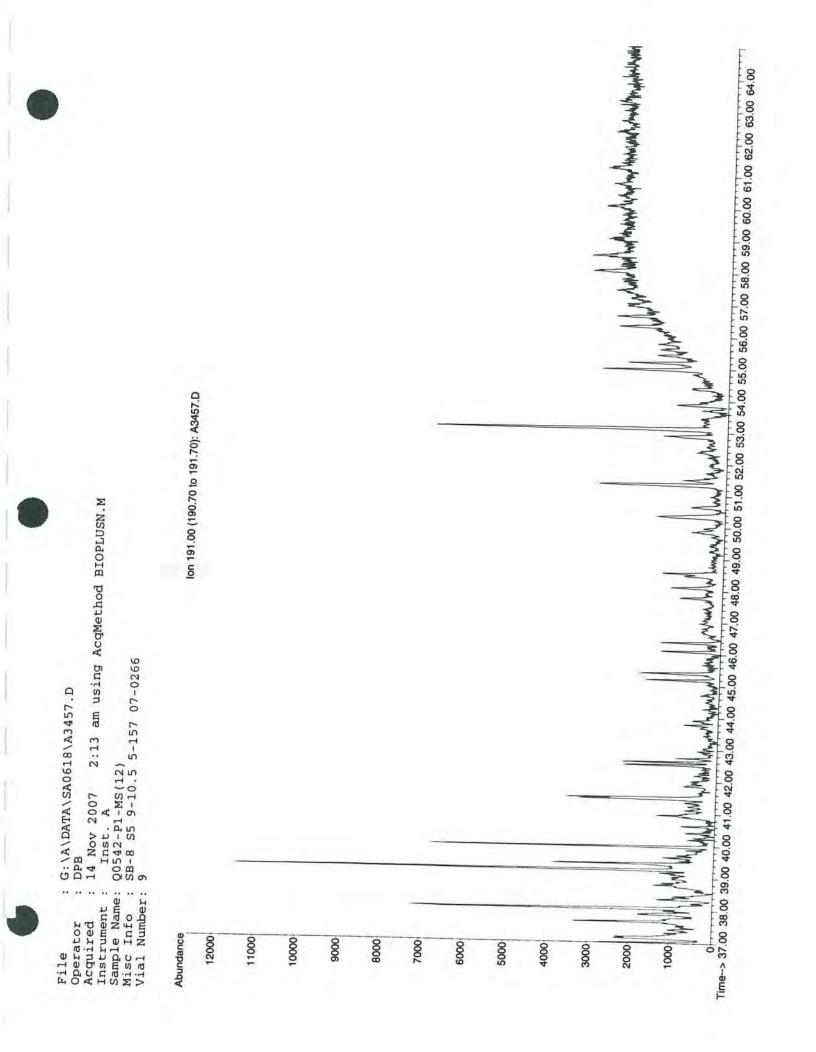
Client ID	GO98: North Slope				
Client ID	Crude				
Battelle ID	BL088NSC-P				
Sample Type	NSC				
Collection Date	10/30/07				
Extraction Date	10/30/07				
Analysis Date					
Analytical Instrument	11/13/07				
% Moisture	MS				
% Lipid	NA				
	NA				
Matrix Sample Size	OIL				
Size Unit-Basis	5.04				
Units	MG_OIL	÷		and the second se	
OTINS	MG/KG_OIL	Target % [Difference	Qualifier	
C23 Tricyclic Terpane	41.31	47.76	13.5		
C29 Tricyclic Terpane -22S	13.23	14.70	10.0		
C29 Tricyclic Terpane -22R	13.32	14.64	9.0		
18a(H)-22,29,30-Trisnomeohopane -TS	15.52	15.96	2.8		
17a(H)-22,29,30-Trisnorhopane -TM	22.68	24.82	8.6		
30-Norhopane	61.75	69.58	11.3		
18a(H) & 18b(H)-Oleananes	01.75 U	09.00	11.3		
Hopane	106.73	120.14	11.2		
30-Homohopane -22S	50.15	59.93			
30-Homohopane -22R	39.69	39.69	16.3		
13b(H),17a(H)-20S-Diacholestane	41.02	44.18	0.0		
13b(H),17a(H)-20R-Diacholestane	24.2	25.52	7.2		
14a(H),17a(H)-20R-methylcholestane	33	33.94	5.2		
14a(H),17a(H)-20S-Ethylcholestane	35.99	35.93	2.8		
14a(H),17a(H)-20R-Ethylcholestane	40.34		0.2		
C21-TAS	18.95	39.17	3.0		
C26-TAS(20S)	14.31				
C26,C27-TAS	53.61				
C27-TAS(20R)	32.3				
C28-TAS(20S)	29.5				
C28-TAS(20R)	31.76				
C21-MAS	6.02				
C22-MAS	2.94				
C27-MAS	6.92				
C27-20R-MAS					
C27-20S-MAS	7.84				
C28-20S-MAS	14.73				
C27-C2920S/R-MAS	9.46				
C29-20S-MAS	3.04				
C29-20R-MAS					
TAS 245	11.36				
MAS_239	U				
MAG_200	U				
Surrogate Recoveries (%)					
Naphthalene-d8	96				
Acenaphthene-d10					
Phenanthrene-d10	94				
Benzo(a)pyrene-d12	97				
5b(H)-Cholane	105				
John - Shoishe	106				

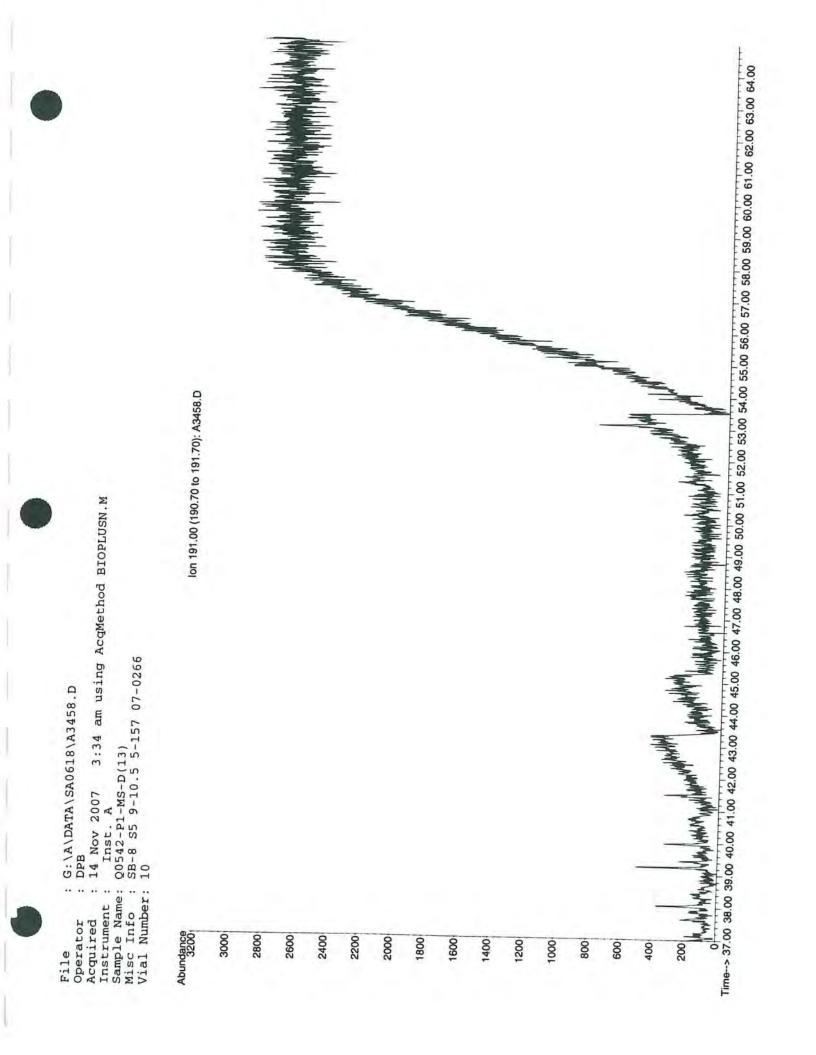




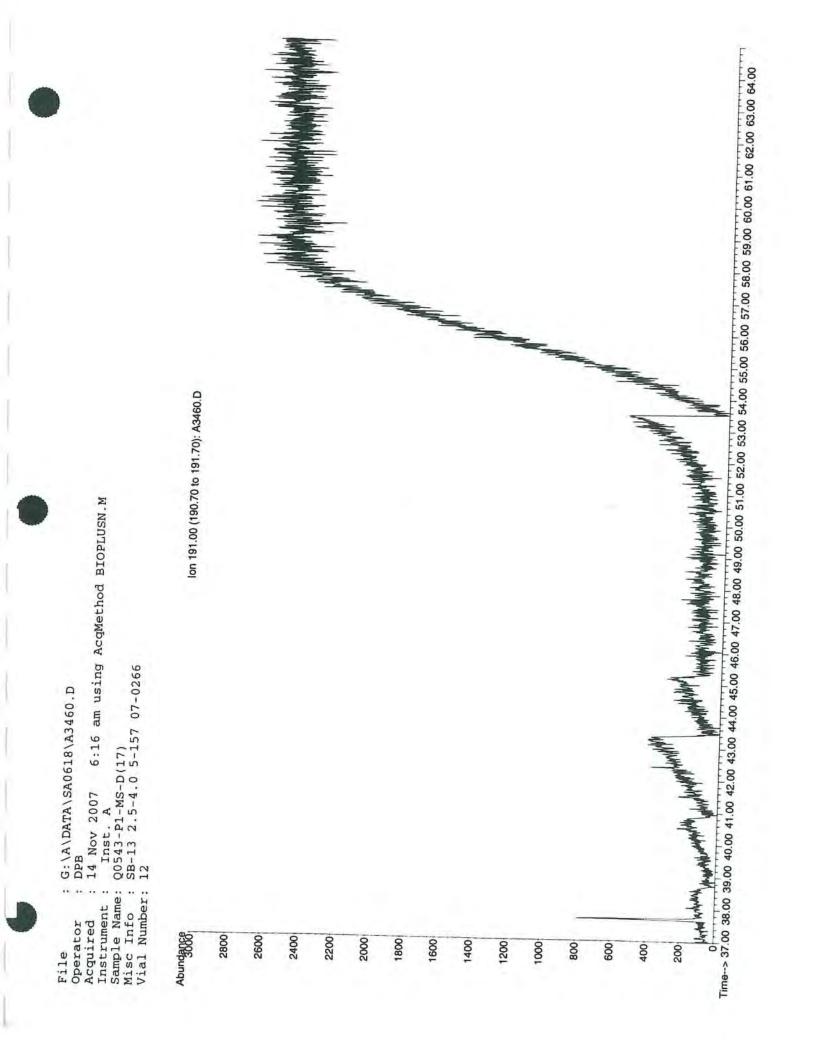




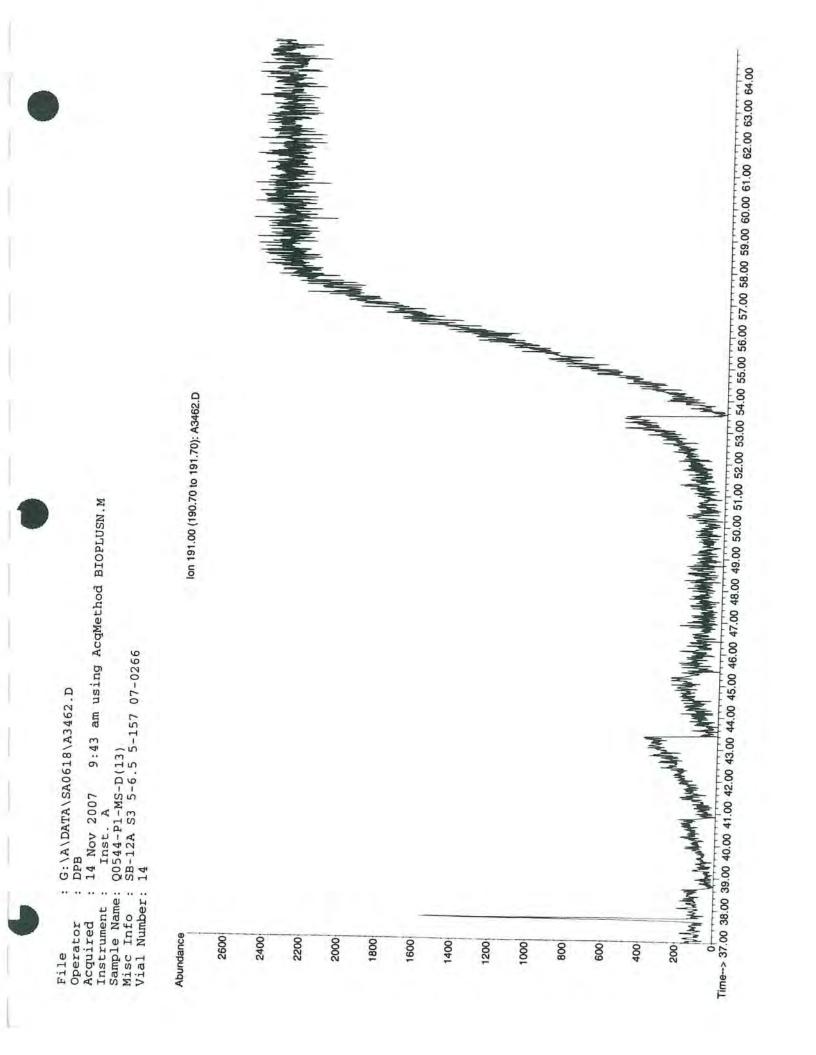


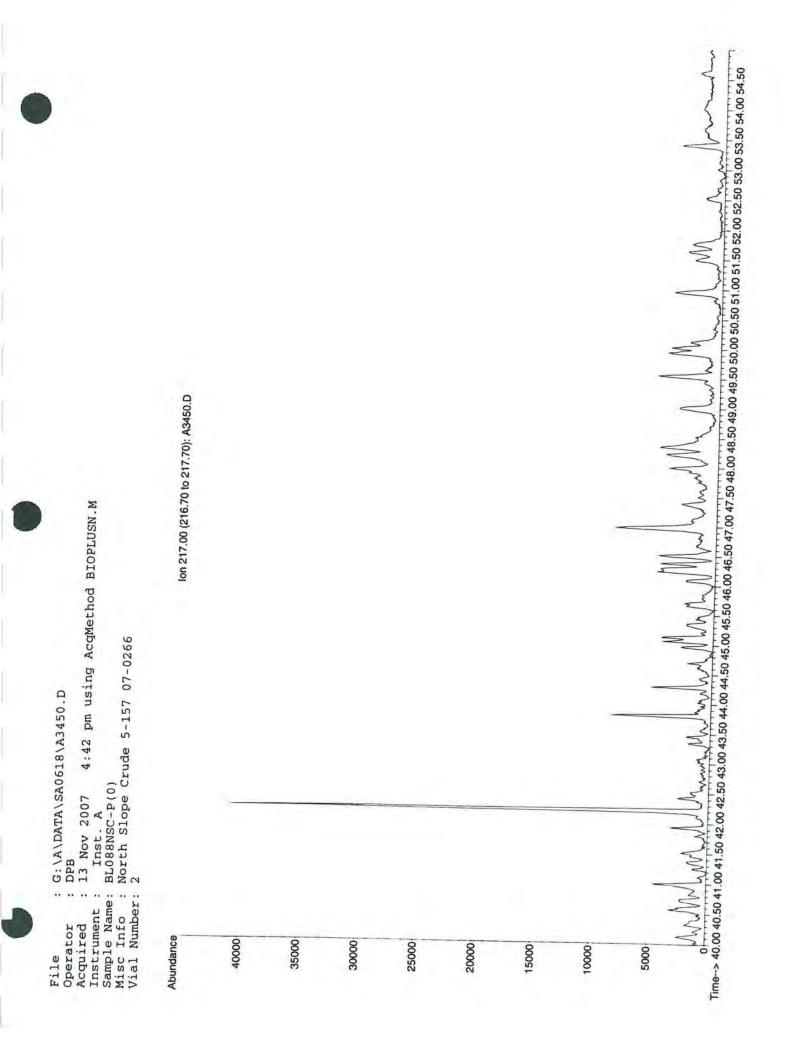


W.	lon 191.00 (190.70 to 191.70): A3459.D									المحمد المحم المحمد	48.00 49.00 50.00 51.00 52.00 53.00 54.00 56.00 56.00 57.00 58.00 59.00 60.00 61.00 62.00 63.00 64.00
: G:\A\DATA\SA0618\A3459.D : DPB : 14 Nov 2007 4:55 am using AcqMethod BIOPLUSN.M : Inst. A : Q0543-P1-MS(16) : SB-13 2.5-4.0 5-157 07-0266 : 11	Ion 191.00 (190.7										0/11-11 10 11 10 11-11 10 11-1
File Operator Acquired Instrument Sample Name: Misc Info	Abundance	20000	16000	0001	12000	10000	8000	6000	4000	2000	me> 37.00 38.00



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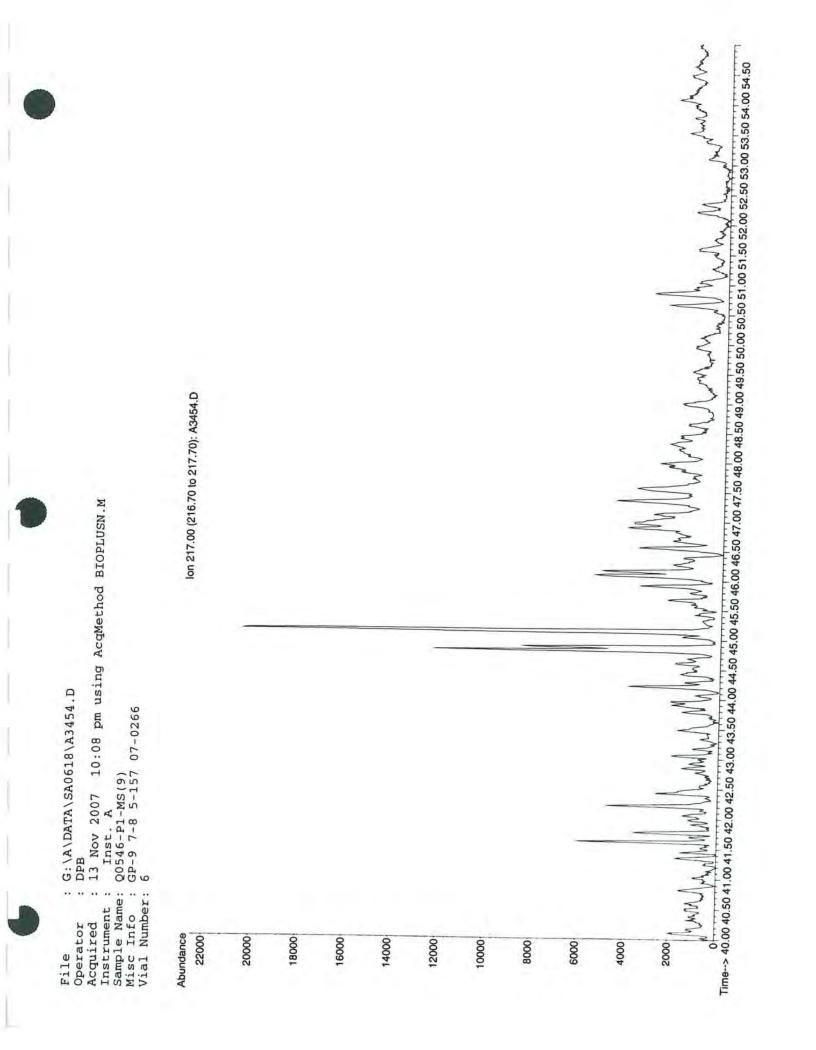


File : G:\A\DATA\SA0618\A3452.D Operator : DPB Acquired : 13 Nov 2007 7:25 pm using AcqMethod BIOPLUSN.M Instrument : Inst. A Sample Name: BL065PB-P-MS(5) Misc Info : Procedural Blank 5-157 07-0266 Vial Number: 4

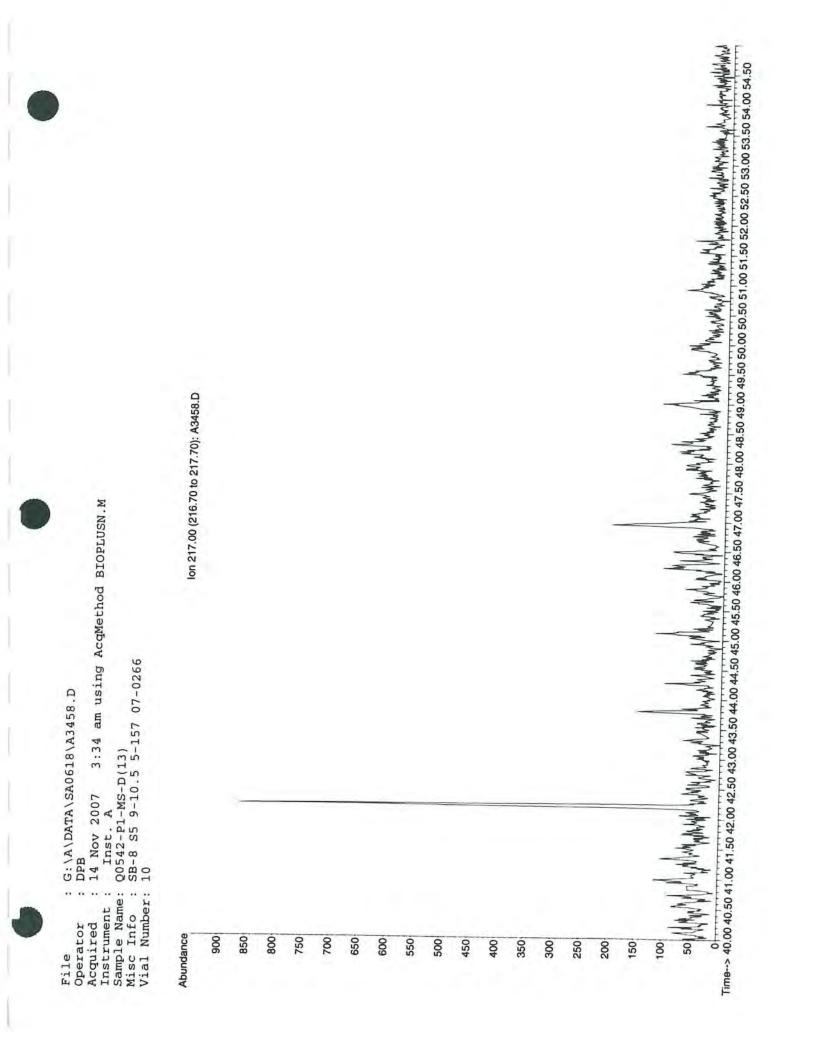
12000	11000	10000	0006	8000	7000	6000	5000	4000	3000	2000	1000
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49.50 50.00 50.50 51.00 51.50 52.00 52.50 53.00 53.50 54.00 54.50 Time

File G:(A)DATA/SA0618\A3453.D Operator Distributions Acquired 11 Bwoy 2007 8:46 pm using AcqMethod BIOPLUSN.M Acquired 11 Bwoy 2007 8:46 pm using AcqMethod BIOPLUSN.M Sample Name: BL066LCS-P-MS(5) Mile Info. 5 Vial Number: 5 Societies Control Sample 5-157 07-0266 MundBQS AbundBQS Societies Control Sample 5-157 07-0266 MundBQS Societies Control Sample 5-157 07-0266 MundBQS Societies Control Sample 5-157 07-0266 MundBQS Societies Control Sample 5-157 07-0266 MundBQS Societies Control Sample 5-157 07-0266 Societies Control	500
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1457.D am using AcqMethod BIOPLUSN.M 7 07-0266	Ion 217.00 (216.70 to 217.70): A3457.D																Mun MMW Lun M	44.00 45.50 45.00 45.50 46.00 46.50 47.00 47.50 48.00 49.50 50.00 50.50 51.00 51.50 52.00 52.50 53.00 53.50 54.00 54.50
<pre>File : G:\A\DATA\SA0618\A3457.D Operator : DPB Acguired : 14 Nov 2007 2:13 am us Instrument : Inst. A Sample Name: Q0542-P1-MS(12) Misc Info : SB-8 S5 9-10.5 5-157 07-(Vial Number: 9</pre>	Abundance 17000	16000	15000	14000	13000	12000	11000	10000	0006	8000	2000	6000	5000	4000	3000	2000	1000 NVVV WWWWWWWWWWWWWWWWWWWWWWWWWWWWWWW	





4:55 am using AcqMethod BIOPLUSN.M Acquired : 14 Nov 2007 4:55 am using Instrument : Inst. A Sample Name: 20543-P1-MS(16) Misc Info : SB-13 2.5-4.0 5-157 07-0266 Vial Number: 11 : G:\A\DATA\SA0618\A3459.D : DPB Operator File

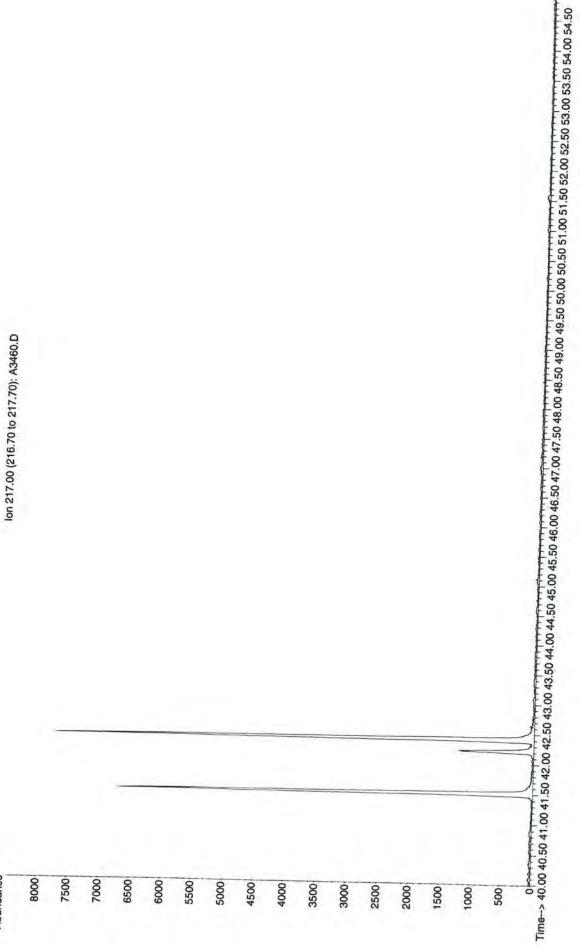
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6:16 am using AcqMethod BIOPLUSN.M Acquired : 14 Nov 2007 6:16 am using Instrument : Inst. A Sample Name: Q0543-P1-MS-D(17) Misc Info : SB-13 2.5-4.0 5-157 07-0266 Vial Number: 12 G:\A\DATA\SA0618\A3460.D DPB Operator Acquired File

Abundance





7:37 am using AcqMethod BIOPLUSN.M Acquired : 14 Nov 2007 7:37 am using A Instrument : Inst. A Sample Name: 20544-P1-MS(12) Misc Info : SB-12A S3 5-6.5 5-157 07-0266 Vial Number: 13 G:\A\DATA\SA0618\A3461.D DPB Operator Acquired Instrument File

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Ion 217.00 (216.70 to 217.70): A3461.D

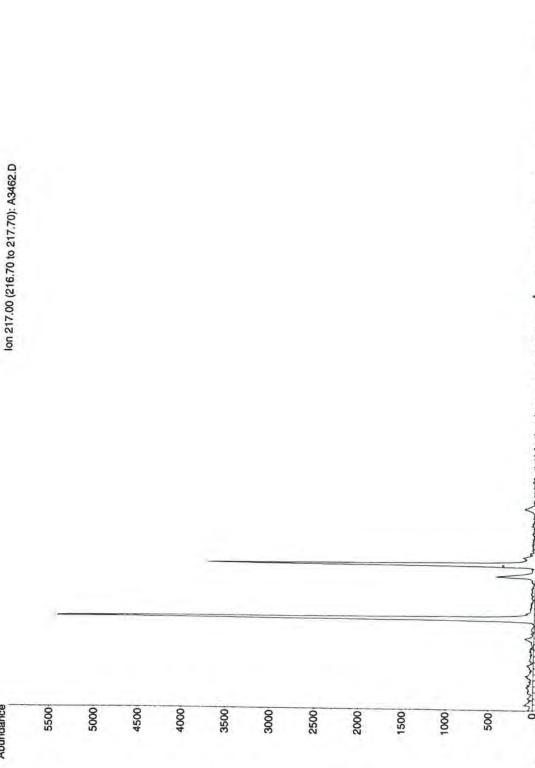


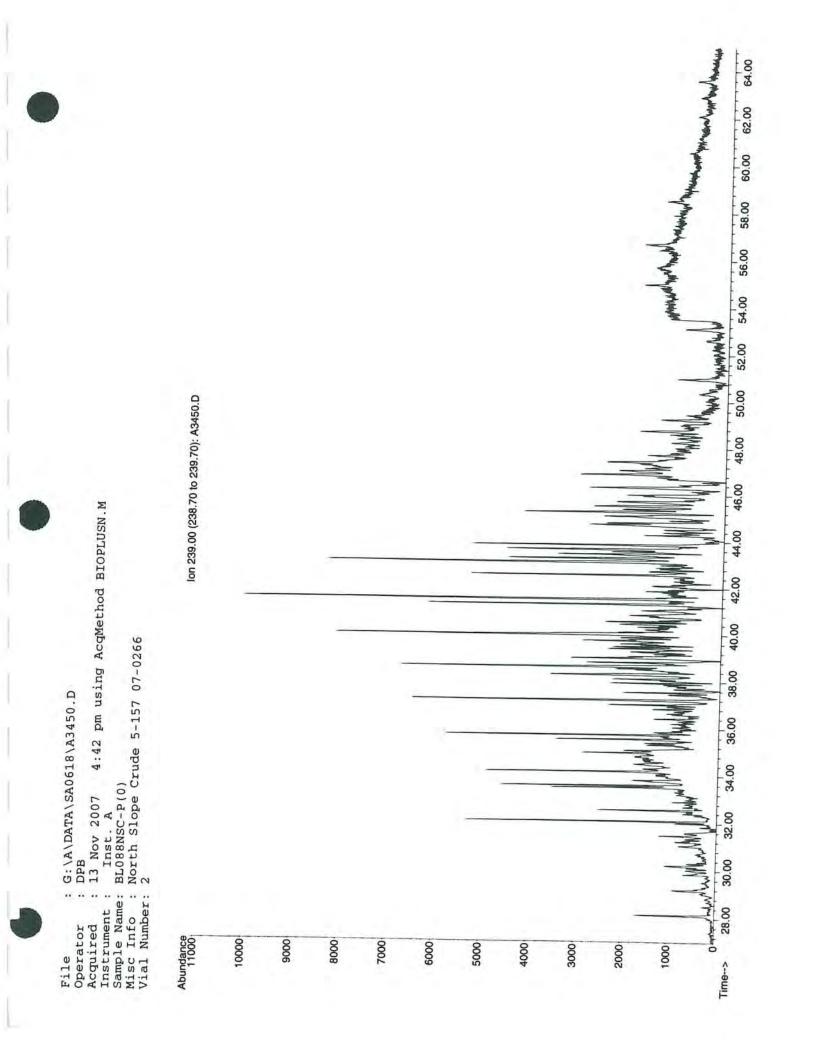
0 0 0 40.50 41.00 41.50 42.00 42.50 43.50 44.00 44.50 45.00 45.50 46.00 45.50 47.00 47.50 48.00 48.50 49.00 49.50 50.00 51.50 51.00 51.50 52.00 52.50 53.50 54.00 54.50



9:43 am using AcqMethod BIOPLUSN.M Instrument : Inst. A
Sample Name: Q0544-P1-MS-D(13)
Misc Info : SB-12A S3 5-6.5 5-157 07-0266
Vial Number: 14 G: \A\DATA\SA0618\A3462.D 14 Nov 2007 DPB Operator Acquired File

Abundance







File : G:\A\DATA\SA0618\A3452.D Operator : DPB Acquired : 13 Nov 2007 7:25 pm using AcqMethod BIOPLUSN.M Instrument : Inst. A Sample Name: BL065PB-P-MS(5) Misc Info : Procedural Blank 5-157 07-0266 Vial Number: 4

64.00

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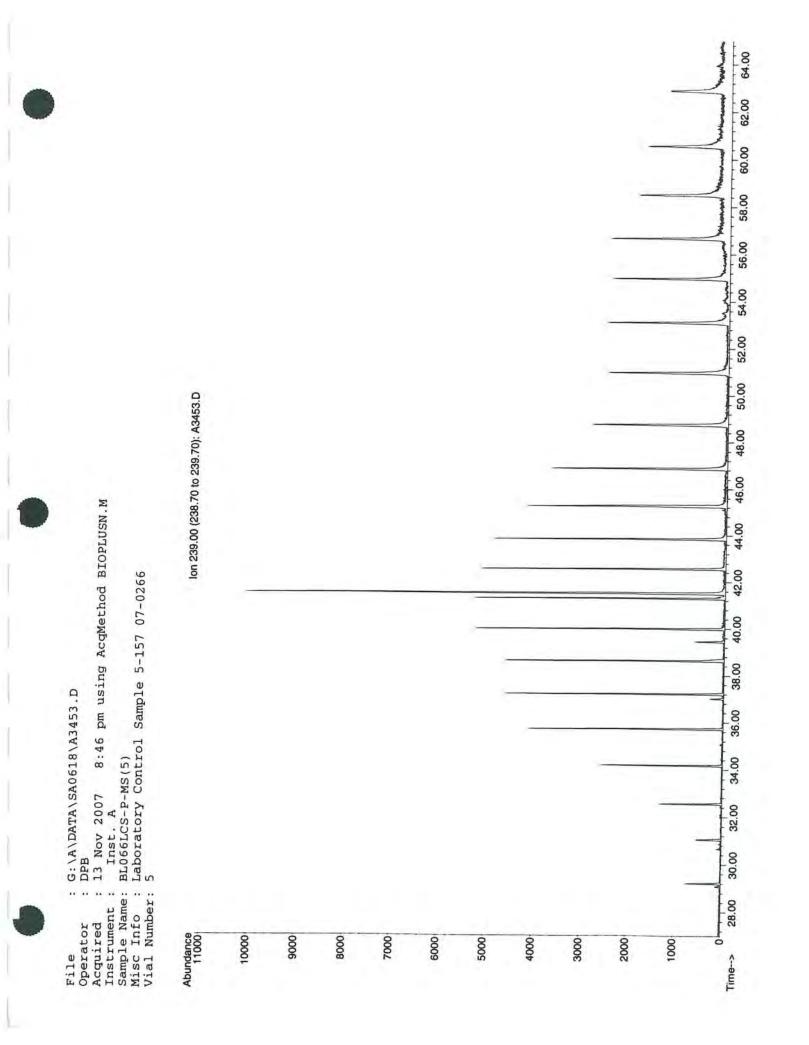
34.00

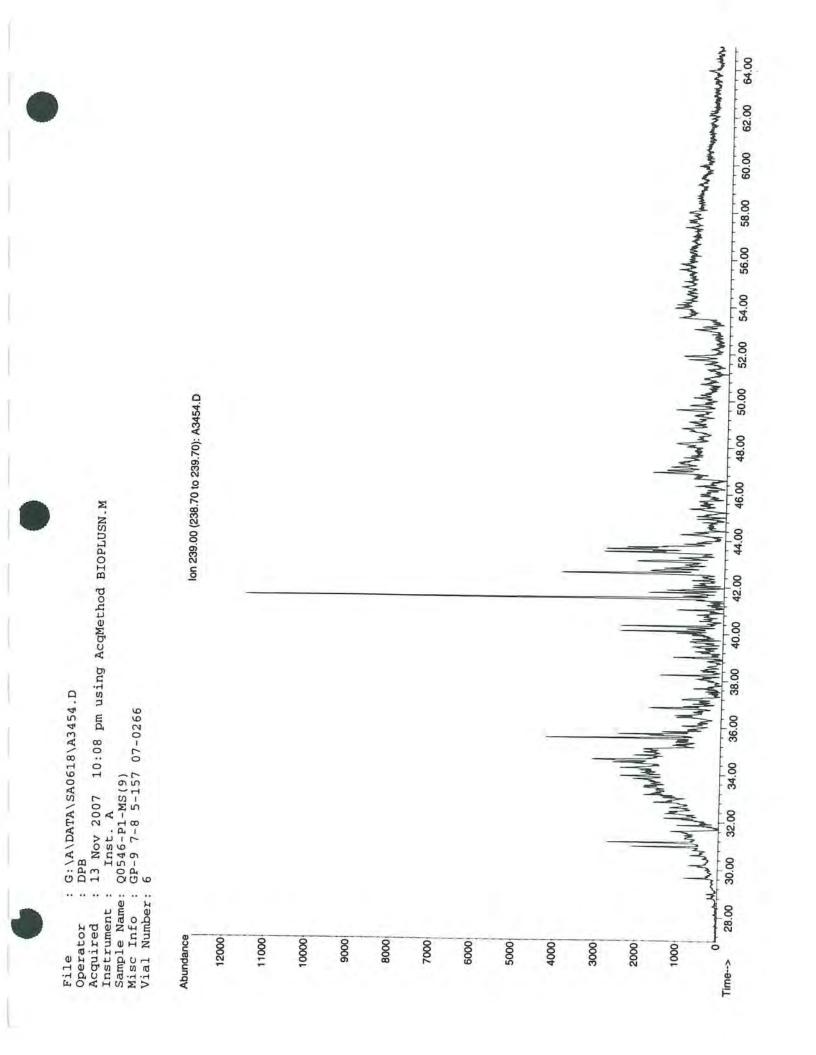
32.00

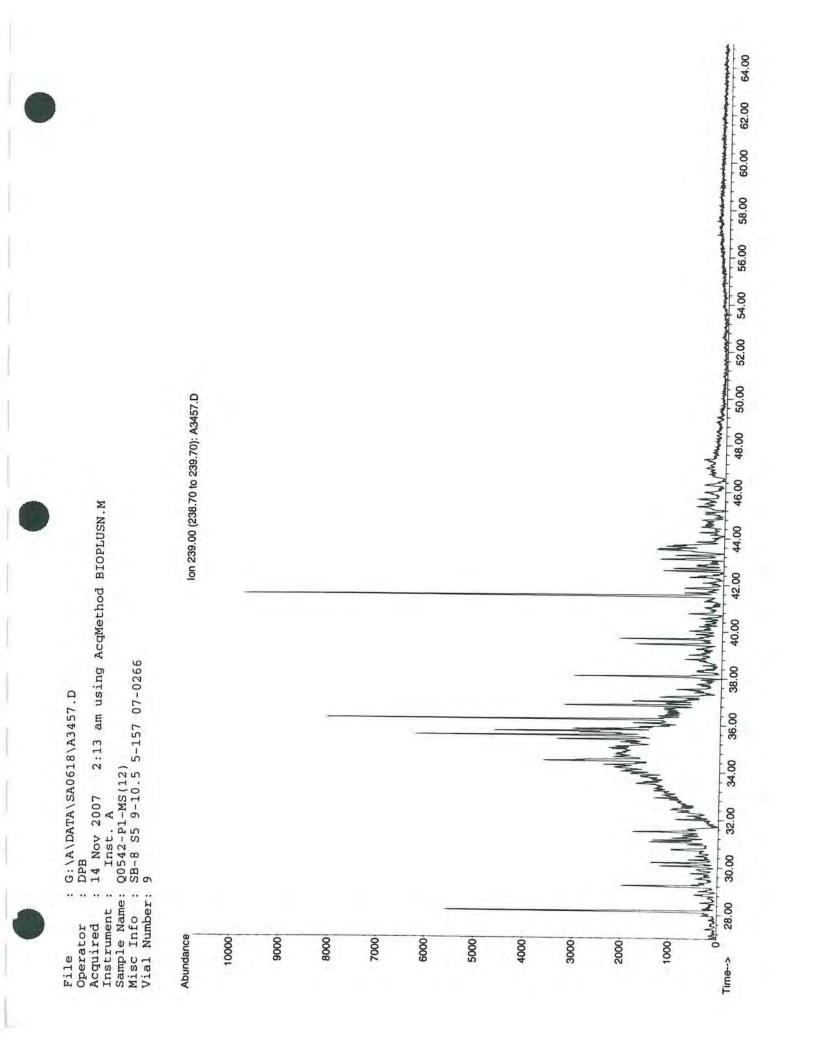
30.00

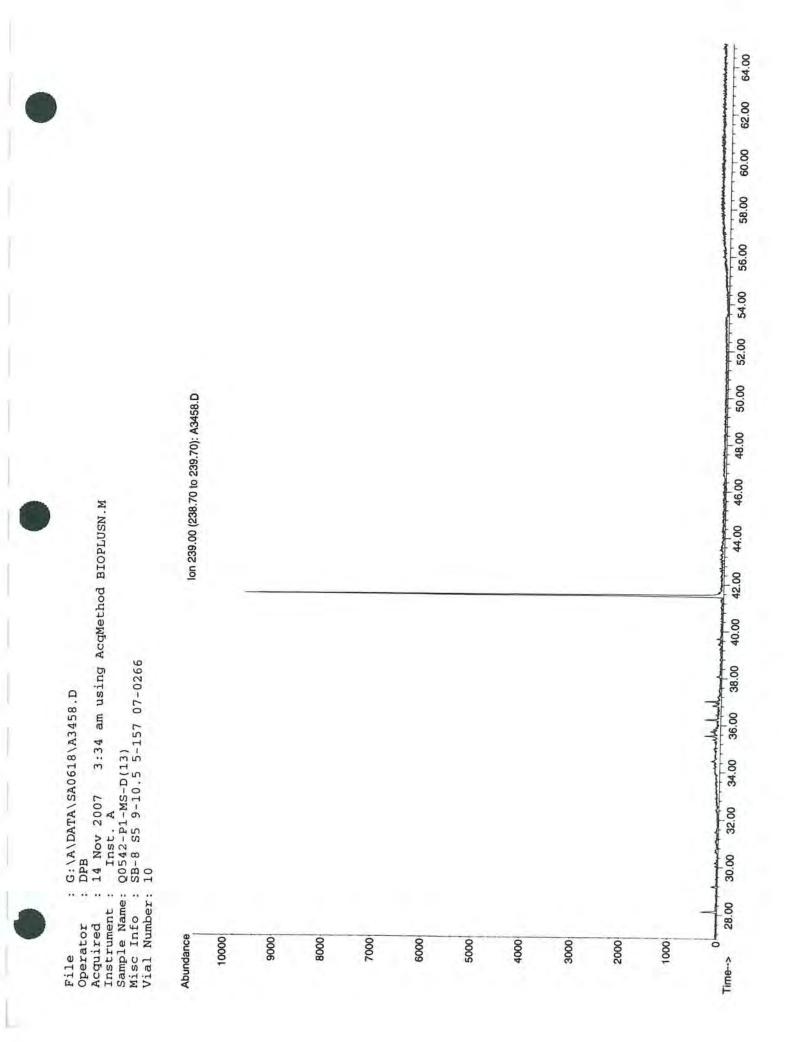
28.00

Time-->

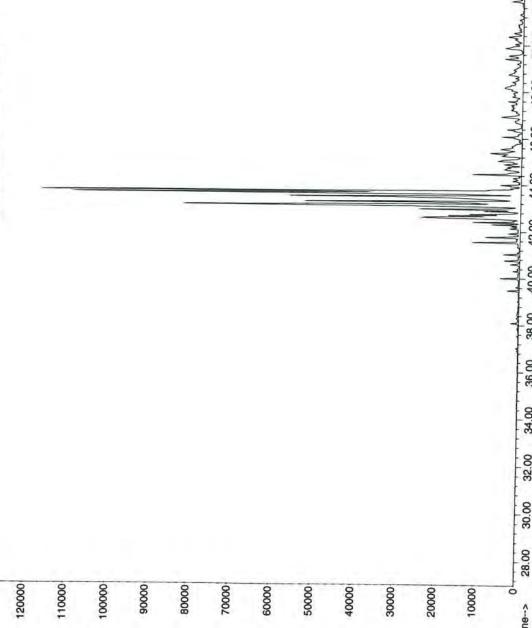




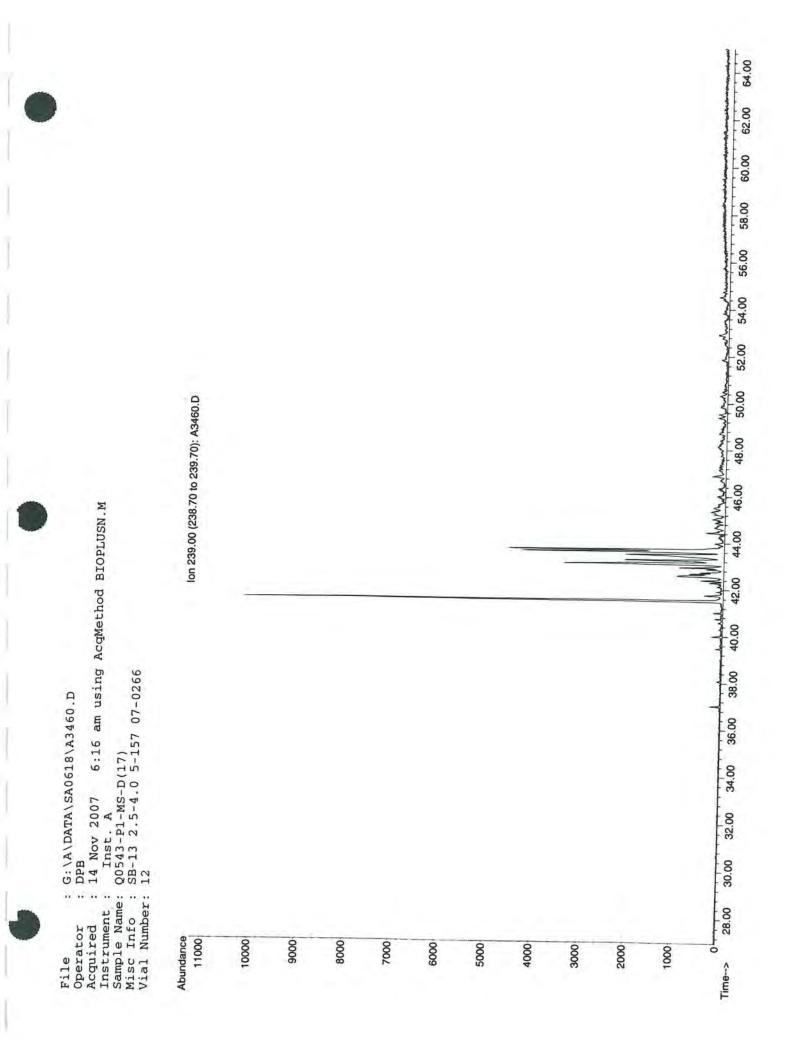




•	File : G:\A\DATA\SA0618\A3459.D Operator : DPB Acquired : 14 Nov 2007 4:55 am using Instrument : Inst. A Sample Name: Q0543-P1-MS(16) Misc Info : SB-13 2.5-4.0 5-157 07-0266 Vial Number: 11	Abundance	120000
	: G:\A\DATA\SA0618\A3459.D : DPB : 14 Nov 2007 4:55 am usi Inst. A : 20543-P1-MS(16) : SB-13 2.5-4.0 5-157 07-02 : 11		
	18\A3459 4:55 am 1 5-157 07 [.]		
	.D ising Acq -0266		
	618\A3459.D 4:55 am using AcqMethod BIOPLUSN.M 6) 5-157 07-0266	lon	
•	OPLUSN.M	Ion 239.00 (238.70 to 239.70): A3459.D	
l :		0 to 239.70):	
		A3459.D	



64.00 62.00 60.00 58.00 56.00 \$ 52.00 54.00 42.00 44.00 46.00 48.00 50.00 52 40.00 38.00 36.00 34.00 32.00 30.00 28.00 Time->





7:37 am using AcqMethod BIOPLUSN.M Operator : DPB Acquired : 14 Nov 2007 7:37 am using Ac Instrument : Inst. A Sample Name: Q0544-P1-MS(12) Misc Info : SB-12A S3 5-6.5 5-157 07-0266 Vial Number: 13 : G:\A\DATA\SA0618\A3461.D : DPB File

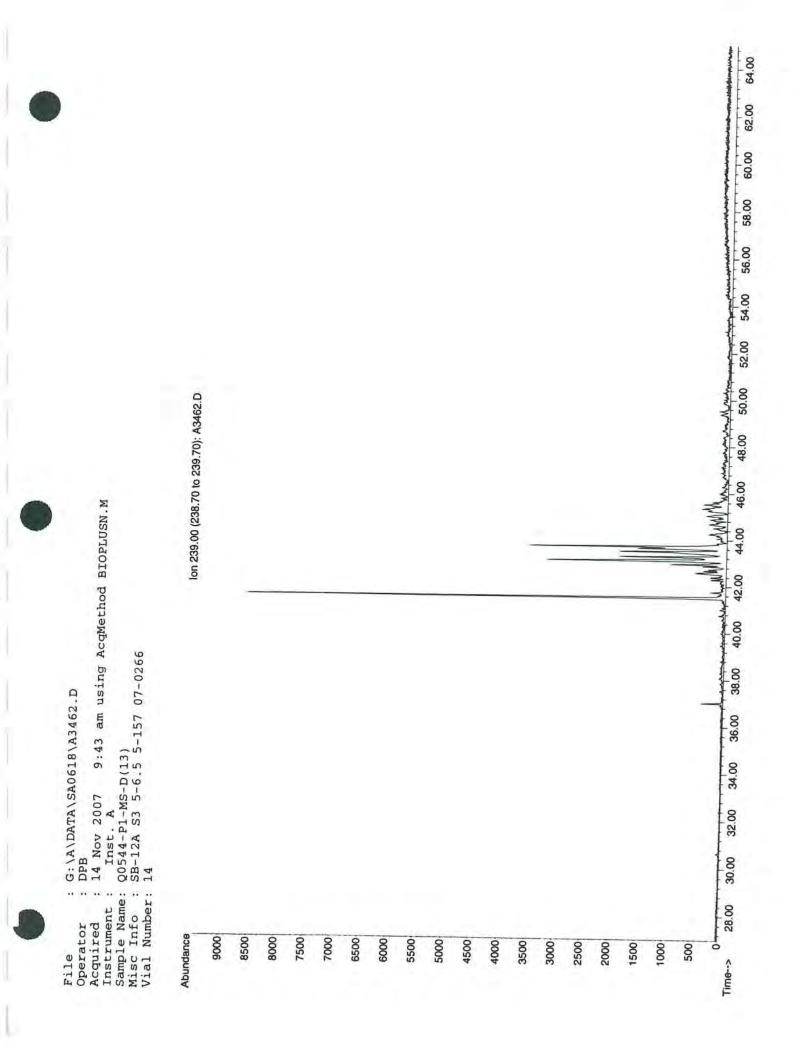
Abundance

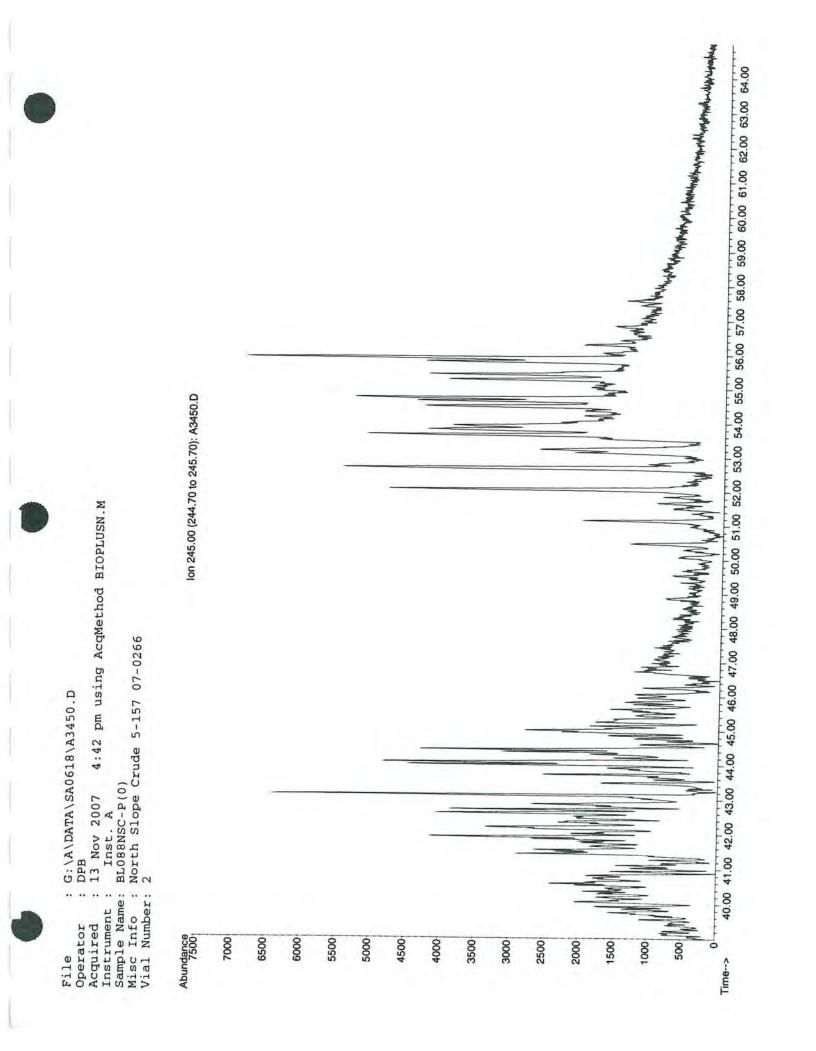
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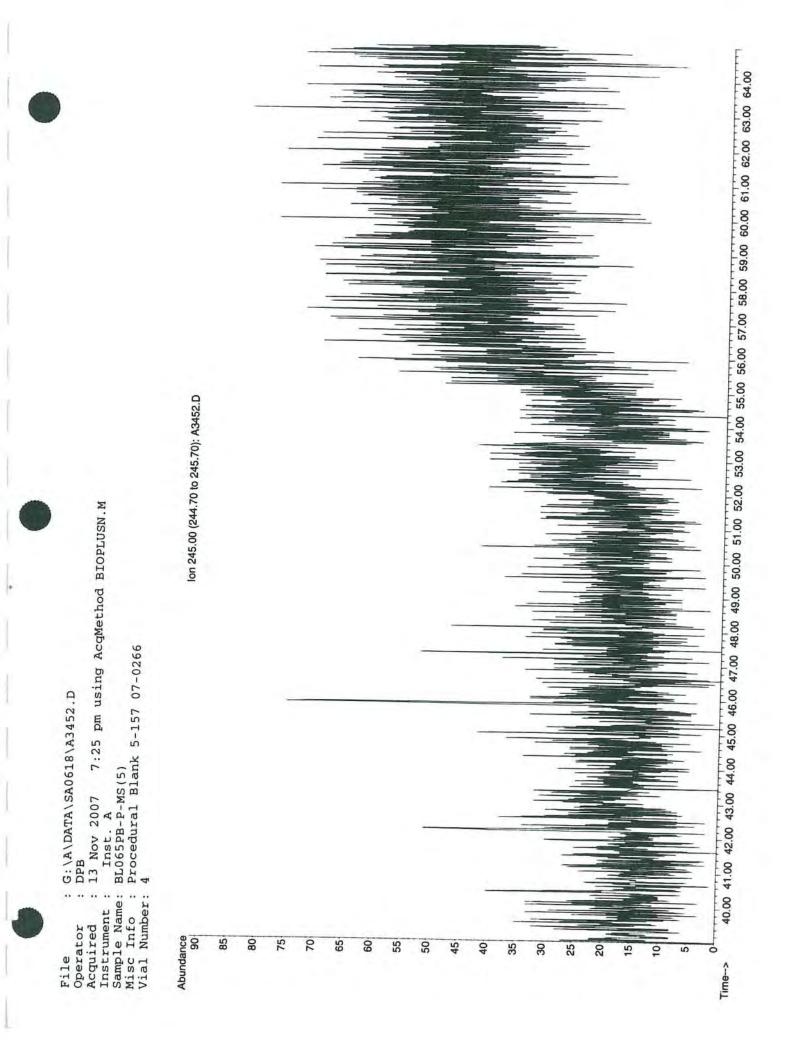
	2	1011 239.00 (238.70 to 239.70): A3461.D	02: 10): Y3401.D		
160000					
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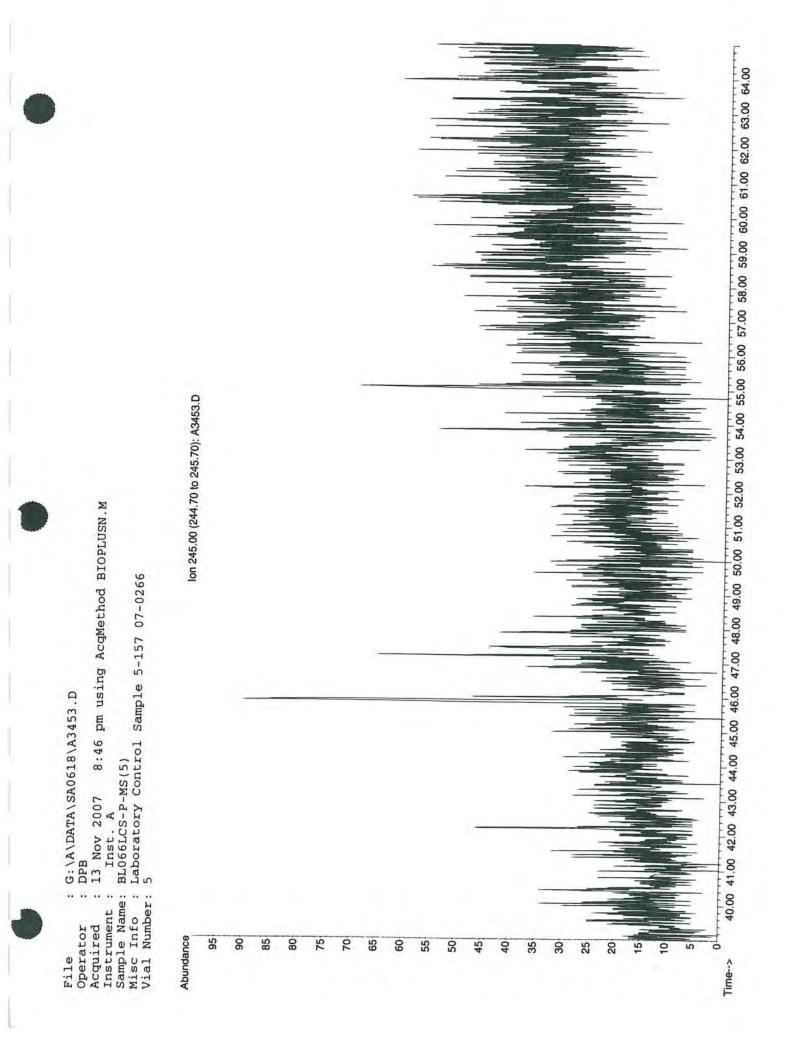
64.00

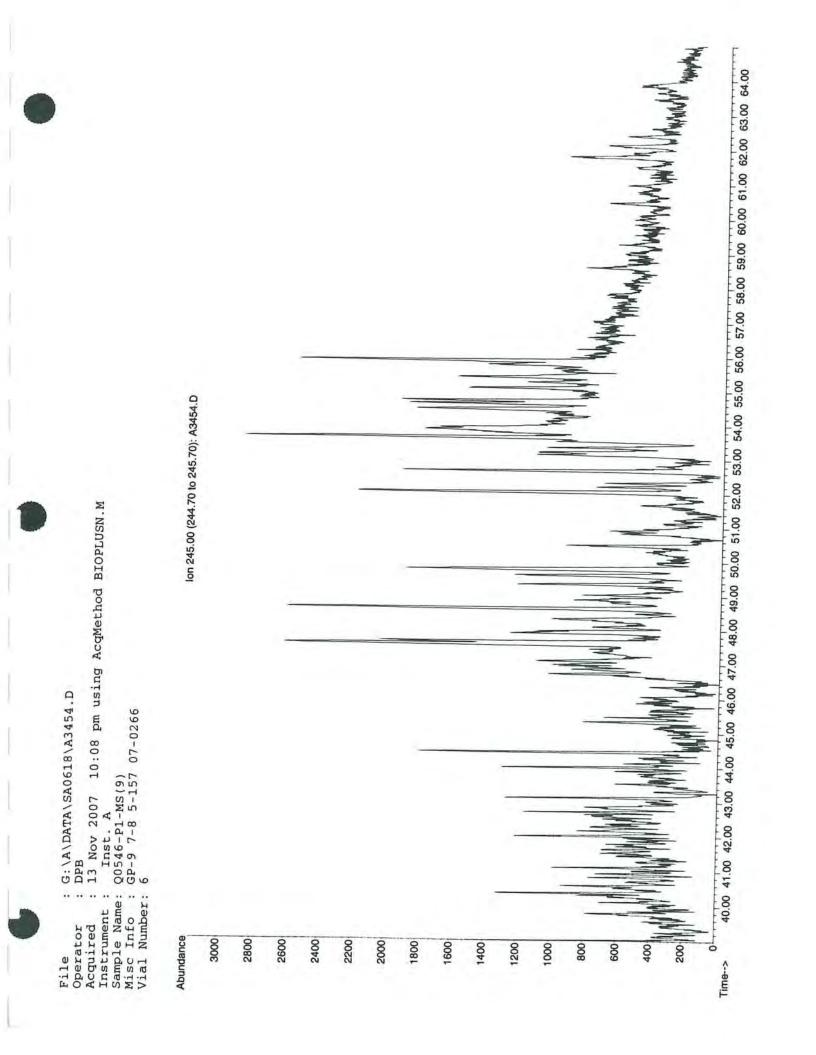
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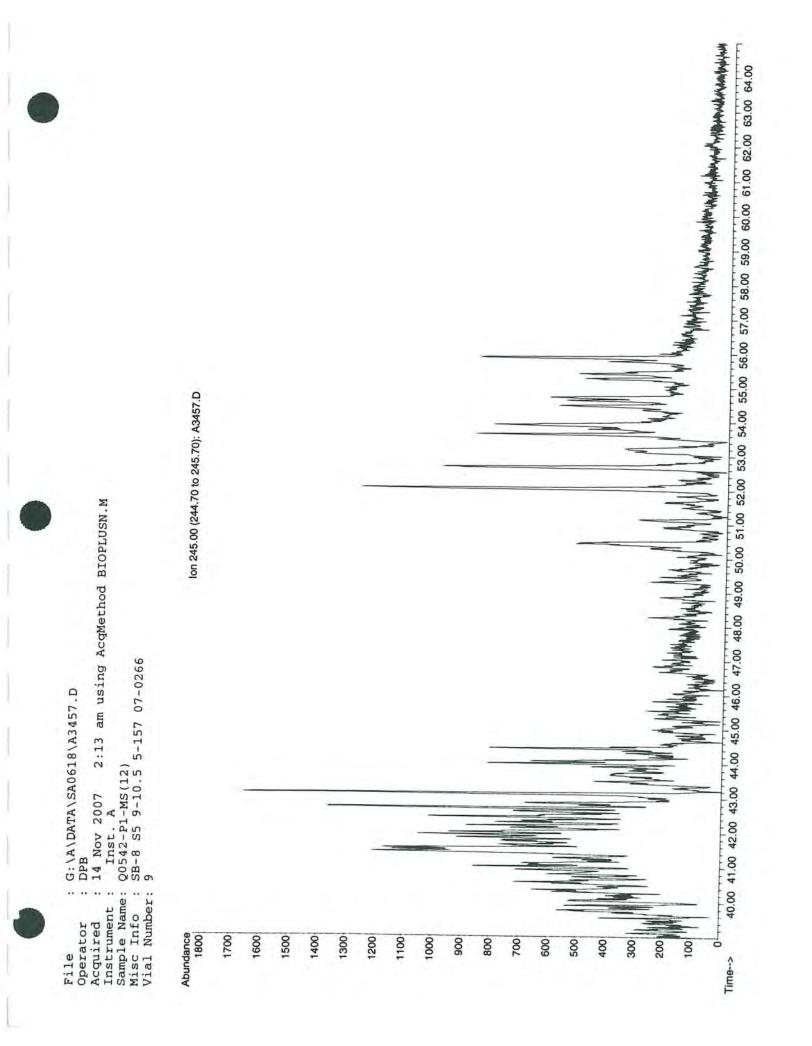


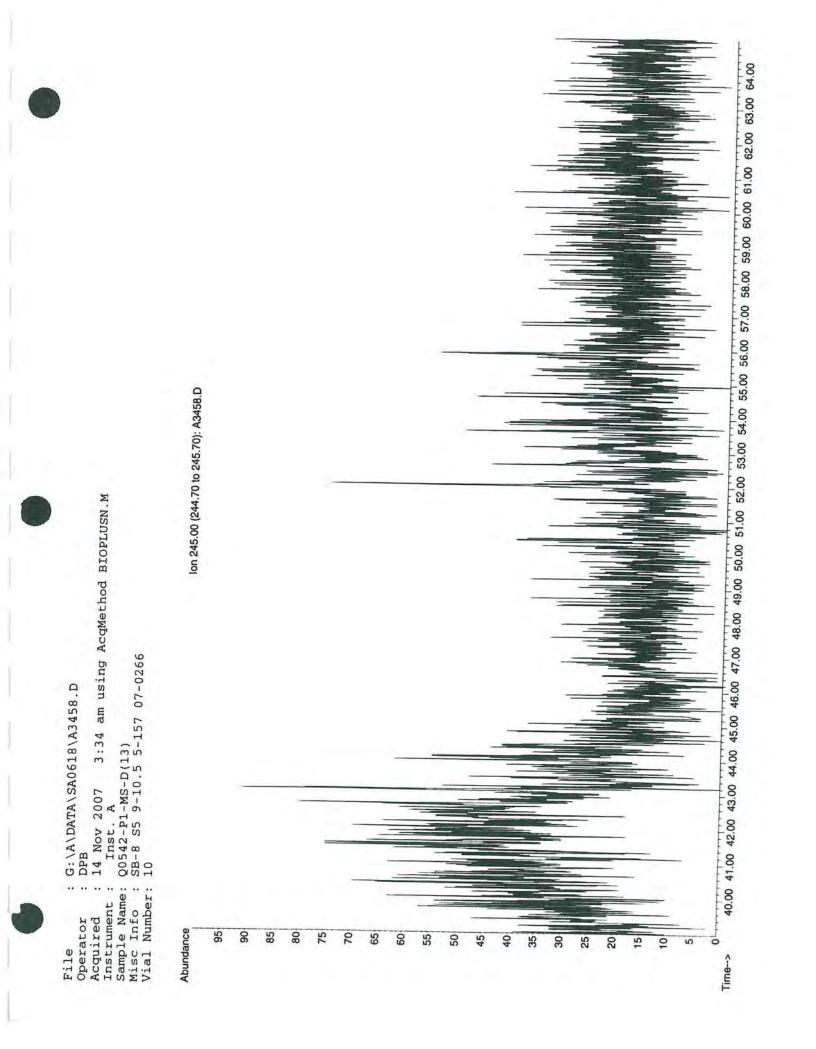


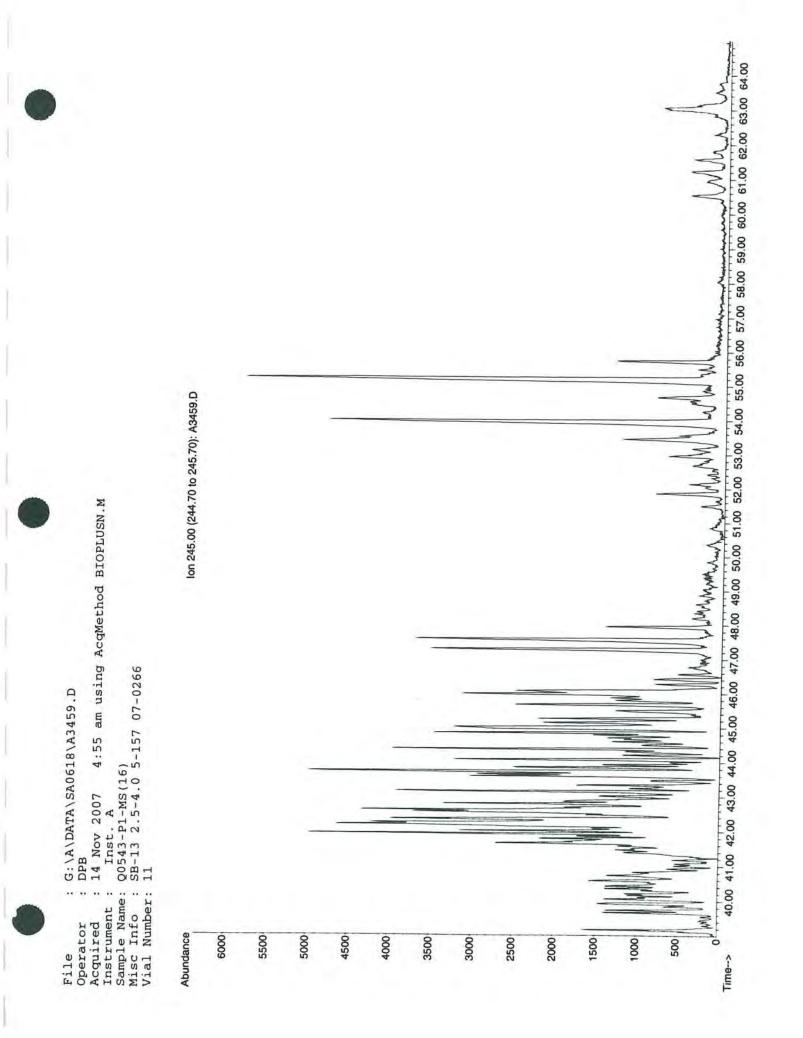


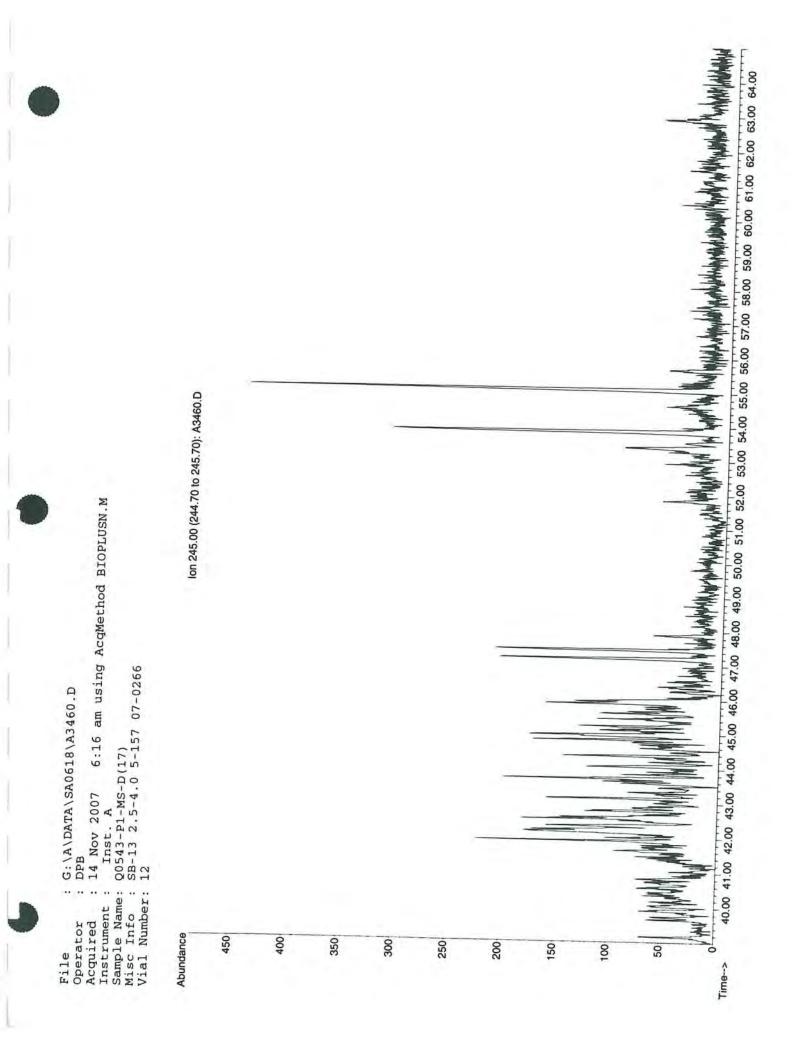


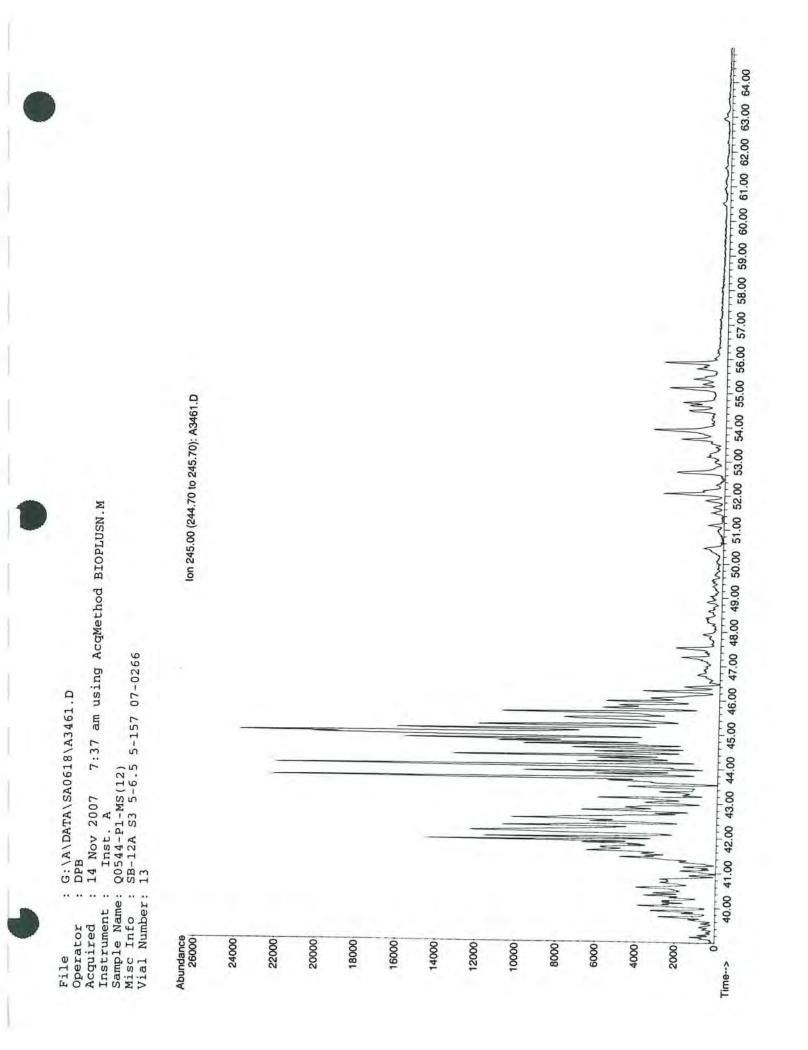


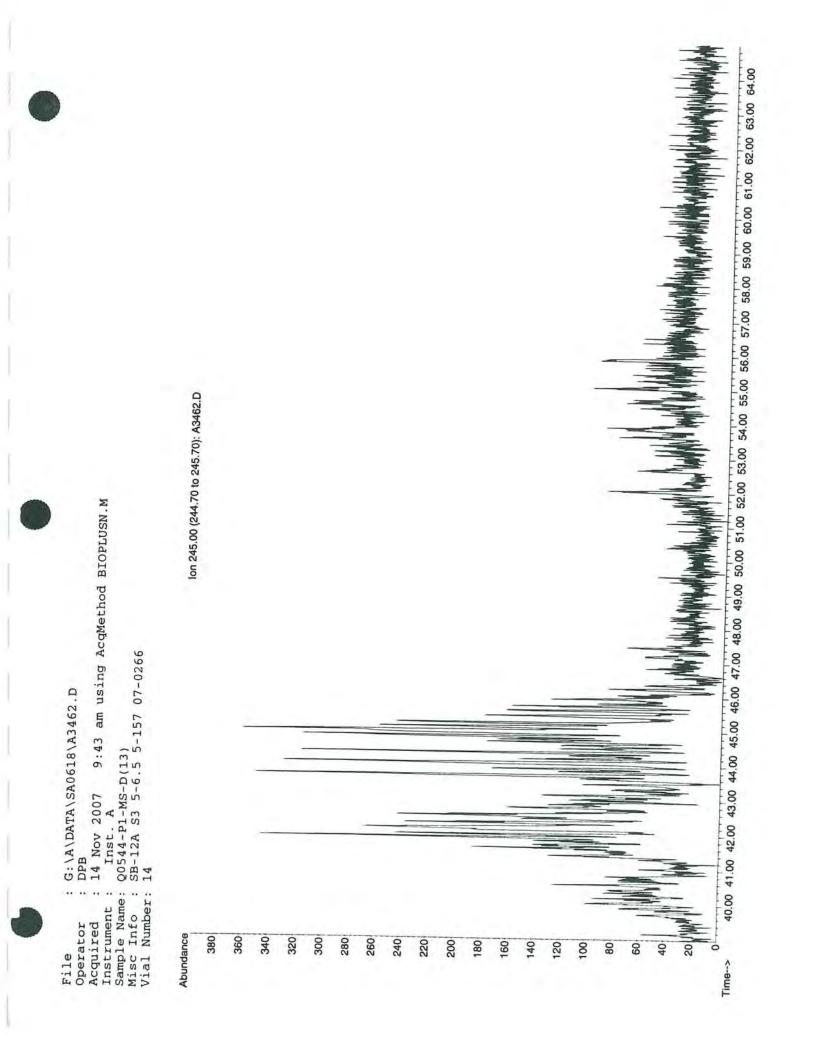












Battelle

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Glossary of Data Qualifiers

Flag: Application:

- B Analyte concentration found in the sample at a concentration <5x the level found in the procedural blank.
- D Dilution Run. Initial run outside linear range of instrument.
- E Estimate, result is greater than the highest concentration level in the calibration.
- H Surrogate diluted out. Used when surrogate recovery is affected by excessive dilution of the sample extract.
- J Analyte detected below the sample-specific Reporting Limit (RL).
- m Confirmation column manually over-ridden by analyst
- ME Significant Matrix Interference Estimated value.
- MI Significant Matrix Interference value could not be determined or estimated.
- n Quality Control (QC) value is outside the accuracy or precision Data Quality Objective (DQO), but meets the contingency criteria.
- N Quality Control (QC) value is outside the accuracy or precision Data Quality Objective (DQO)
- NA Not applicable
- p Dual column value exceeds RPD criteria
- T Holding Time (HT) exceeded.
- U Analyte not detected at 3:1 signal:noise ratio.

ATTACHMENT 2D-7 Stanford Aqueous Equilibrium Report

Final Report – February 3, 2006

PAH Partitioning in Black Carbon-impacted Sediments from Lake Union

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SUMMARY

This report summarizes laboratory investigations of the physicochemical nature of polycyclic aromatic hydrocarbon (PAH) contamination in sediment samples from Lake Union, Seattle, WA. This work infers the extent to which black carbon (BC) materials are present in samples, and the role that these black, particulate organic particles may have in affecting the PAH availability from sediments. Three partitioning models are evaluated to explain the aqueous equilibrium sorption between PAHs and sediment samples: a conventional partitioning model based on natural organic matter sorption, an oil or tar phase partitioning model based on Raoult's Law assuming an ideal aromatic phase, and a model assuming sorption on soot-like matter.

Twenty sediment samples were examined in this study, which exhibited a wide variety of physicochemical characteristics. It was noted that one sample, CR10-NAPL, appeared completely different from the other nineteen sediment samples and was comprised mainly of a soft-tar like non-aqueous phase liquid (NAPL) substance. The total organic carbon contents of these samples ranged from less than 1% to over 70% by weight, and PAH analyses showed highly variable degrees of PAH contamination in these sediment samples. The total concentrations of the 16 EPA priority PAHs ranged from 18 mg/kg for NLU 54 to 4,400 mg/kg for NLU 51, and for the NAPL sample the value was 90,500 mg/kg. In addition to PAH analyses in bulk sediment samples, PAH analyses on density-separated fractions provided insights about the association of PAHs with sediment particles. The significant findings were that PAH concentrations in the low-density fractions; and for most samples the majority of the PAH mass (78 ~ 100% of total PAH mass) was associated with the low-density particulate fractions. These observations were not surprising given the hydrophobicity of PAH compounds and the high organic matter and

black carbon content in the low-density fractions.

Measurements of aqueous equilibrium concentrations of PAHs from these sediment samples provide a quantitative evaluation of PAH leaching potential and availability from the sediment to the aqueous phase. Site-specific, sediment-water distribution coefficients (K_d) of PAHs are calculated. By comparing experimentally-determined K_d values with predictions based on different partitioning models, we propose three PAH partitioning scenarios occur in these sediment samples: *i*) modified fraction organic carbon partitioning: the majority (fifteen out of twenty in total) of sediment samples generally agree with the conventional organic carbon equilibrium partitioning model, $K_d = f_{oc} K_{oc}$. However, due to the existence of black carbonaceous particles, the observed K_d values are approximately 0.5 ~ 1.0 logarithmic units higher than conventional predictions for sorption onto natural organic matter with the divergence from the conventional model greater for the more hydrophobic PAH compounds; ii) oil / tar partitioning: three sediment samples NLU 51, NLU 56, NLU 65, and the NAPL sample (CR10-NAPL) exhibit PAH partitioning behavior described by an oil/tar-water partitioning model, which includes the case of oily-coated soot. This model assumes that the PAHs are present in an oil phase and that the partitioning behavior follows the dissolution process from the oil phase; and iii) soot carbon partitioning: NLU 73-Stanford is the only sample that clearly exhibits partitioning behavior of PAHs being controlled by oil soot (or lampblack, carbon black). In this case the PAH sediment-water distribution coefficients K_d are generally 1.5 ~ 2 orders of magnitude higher than those derived on the basis of conventional natural organic matter partitioning, and correspondingly the aqueous equilibrium concentrations are disproportionally lower than predicted by such partitioning models.

This report also includes two documents as appendixes. Appendix I is the detailed data for PAH analyses on density separated fractions for each sediment sample. Appendix II comprises results on aqueous equilibrium concentration measurements and calculated site-specific PAH sediment-water distribution coefficients for sediment samples.

INTRODUCTION

This project examined the characteristics of polycyclic aromatic hydrocarbon (PAH)

contamination in sediment samples from Lake Union, Seattle, WA, and the role that black carbonaceous particles may contribute to the presence of PAHs and PAH partitioning behaviors to the aqueous phase. Specifically, this project (1) inferred the extent to which the sediment samples from Lake Union contain black carbon materials, and (2) investigated their roles in affecting the availability of PAHs in samples obtained from the site.

This project evaluated the sorptive binding of PAHs to the lake sediment in order to better determine how sorption affects the fate of these contaminants in the aquatic environment. It is well accepted that hydrophobic organic compounds such as PAHs sorb to the organic carbon fraction of sediment. Thus, sorption models based on the fraction organic carbon content (f_{oc}) are useful to predict the sorption behavior of hydrophobic organic compounds to natural sediments assuming the organic matter to be homogeneous and equally sorptive (1, 2). Traditionally such models employ an organic-carbon-normalized distribution coefficient (K_{oc}) that is dependent on the sorbing chemical's hydrophobicity or solubility, as may be quantified by the sorbate's octonal-water partition coefficient K_{ow} (3). Although the organic carbon fraction foc is generally taken as a measure to characterize the sorption capacity of natural sediments, investigators have noted for several years that the sorption of hydrophobic organic compounds to sediments is not always consistent with model predictions. Our earlier work (4, 5) demonstrated that organic carbon in sediments may comprise different compositional forms having very different sorption capacities. Thus, the nature of the organic carbon in the sediment could have a significant impact on the sorption behavior of contaminants associated with the sediment. This is especially the case if the sediment contains black carbon particulate matter known to have high sorption capacities and strong affinities for PAHs. The strong binding between black carbon materials and PAHs could result in less availability of PAHs to the aqueous phase and thus considerably reduce the potential environmental risk to the aquatic environment.

Furthermore, in our recent work we demonstrate that the presence of oil/tar in soil or sediment also could be an important factor in determining the organic contaminant partitioning behavior of PAHs (5) or PCBs (14). We showed for lampblack that, when the oil/tar content in soil or sediment is high enough to exhaust the sorption capacity of lampblack carbonaceous matter, the oil/tar phase governs PAH partitioning and that PAHs associated with a liquid or

semi-soft oil/tar phase will be more available. This could result in higher aqueous equilibrium concentrations and pose greater environmental concern than PAHs sorbed only on lampblack.

OBJECTIVES AND SCOPE OF THE STUDY

The primary objective of this research is to identify the extent of black carbon materials in Lake Union, WA, sediment samples and to characterize the association of PAHs with black carbon-impacted sediment. The second major objective of this project is to measure aqueous equilibrium concentrations with sediment samples and to calculate site-specific PAH partition coefficients between sediment samples and water. The measured PAH partition coefficients are compared with conventionally-estimated partition coefficients based on the organic carbon content in the sediment. This comparison will shed light on estimating the impact of black carbonaceous materials in the sediment and the effects of such materials on the availability of PAHs from the site.

This report summarizes results of (1) characterization of PAH concentration levels and distribution patterns, as well as (2) the determination of PAH aqueous equilibrium concentrations for sediment samples from Lake Union. We received 16 sediment samples on January 20, 2005, and four remaining sediment samples arrived on February 3, 2005. Since receipt, the following tasks have been conducted on these twenty sediment samples:

1. Characterization of sediment samples and PAH analyses

- A density separation technique aided by a saturated cesium chloride solution was used to identify the percentage by weight light carbonaceous particulate fraction in each sediment sample.
- The total organic carbon content in each sediment sample was measured using a combustion technique followed by nondispersive infrared detection.
- The percentage by weight oil/grease in the whole samples was determined by following procedures similar to SW-846 EPA method 9071A (Hexane Extractable Materials) for oil and grease in soils and sediments using ultrasonic extraction.
- Analyses for 16 EPA priority PAHs were performed on the whole samples and the separated fractions by following EPA method 3550B (ultrasonic extraction), EPA

method 3630C (silica gel cleanup), and EPA method 8100 for PAH analyses with a gas chromatograph (GC) and a flamed ionization detector (FID).

- 2. Aqueous equilibrium tests with sediment samples and calculation of site-specific partition coefficients of PAHs
 - Aqueous equilibrium tests with sediment samples were performed using a newly-developed polyoxymethylene (POM) solid phase extraction (SPE) method as described in Jonker et al., 2001 (6). This technique has been fully characterized with the polyoxymethylene material in our laboratory (7) and is confirmed to be comparable with other aqueous equilibrium concentration measurement techniques such as air-bridge or alum flocculation procedures.
 - The PAH aqueous equilibrium concentrations for selected samples were measured by employing a direct contact protocol, which involves supernatant treatment by alum flocculation and centrifugation. This was performed on six sediment samples to confirm results obtained by the POM solid phase extraction method.
 - An air-bridge technique was used to measure PAH aqueous equilibrium concentrations for the non-aqueous phase liquid sample (CR10-NAPL-1523). This technique was chosen due to the need for absolute certainty of eliminating the potential oil contamination in the aqueous phase measurements.
 - Site-specific partition coefficients (K_d) of PAHs were calculated for each sediment sample, and these results were compared with values obtained based on a conventional, organic-carbon partitioning model. Organic-carbon normalized partition coefficients (K_{oc}) were also determined.

SITE DESCRIPTION AND SAMPLE INFORMATION

The study area of this project comprises a band of sediments in the region surrounding the Gas Works Park site in north-central Lake Union. The Gas Works Park site is located on the northern shore of Lake Union in Seattle, WA. Historically significant activities occurring at this site over time included: a manufactured gas plant, a tar refinery, and park construction. The gas works plant was built on the site in 1906 and the plant gasified coal to produce "town gas." In 1937, oil-gas generators were added and the plant was converted to an oil gasification process.

The gas making operations ceased in 1956 due to the availability of cheaper natural gas. A tar refinery was operated on the northwestern part of the site from 1907 through the 1950s. The tar company processed tars from the gas works plant, as well as feedstock from other sources, to produce various grades of tar and pitch. After the gas works converted to oil gas in 1937, manufactured gas plant tar was no longer a major feedstock to the tar refinery. The property was purchased by the City of Seattle in 1962 and converted to a public park, named Gas Work Park, and opened in 1976. Sediments in Lake Union are very likely contaminated from a number of sources owing to the long history of industrial and commercial activities near or along the lake.

EXPERIMENTAL METHODS

Twenty sediment samples from Lake Union were shipped to our laboratory in ice-packed coolers, and stored at 4 °C until use. Prior to experiments, all sediment samples were wet sieved to remove particles larger than 2 mm in diameter. These large particles were primarily comprised of stones and brick fragments.

Chemicals and Materials. Pesticide grade solvents hexane, acetone, pentane, cyclohexane, and methylene chloride were purchased from Fisher Scientific (Pittsburgh, PA). Solvents were checked regularly for any PAH contaminations by GC-FID as described below. Anhydrous sodium sulfate (Fisher Scientific) was prepared with drying in an oven at 105 °C for 24 hours prior to use. Silica gel (Fisher Scientific) used in the column cleanup procedure was activated at 130 °C for 16 hours. White crystalline powder cesium chloride (CsCl) with a purity > 98% was obtained from Sigma Aldrich, Inc. PAH surrogate solution pyrene- d_{10} (500 ppm in acetone), internal standard 2-fluorobiphenyl, and standard solutions of 16 EPA PAHs for GC calibration were purchased from ULTRA Scientific, Inc. (North Kingstown, RI). Polyoxymethylene (POM) (trade name: poly-acetal, also known as acetal; molecular formula: [-OCH₂]_n; density: 1.38 g/cm³) was obtained from Vink NV, Industriepark 7, B-2220 Heist-op-den-Berg, Belgium. The POM sheet has a thickness of 0.58 mm and natural white color. It was cut into strips with desirable dimensions of 13 mm by 5 mm prior to use and pre-cleaned by soaking in methanol and hexane sequentially for 30 minutes each, then rinsed with Milli-Q[®] water and allowed to air-dry. Before employed in experiments, several strips were randomly selected as blank samples to go through

the whole procedure from extraction, column cleanup, and GC analysis to ensure that no contamination exists with these POM strips.

Density Separation. A saturated cesium chloride solution with a specific gravity of 1.8 was used to separate lighter-density carbonaceous particles from the mineral fraction of soil (i.e., sand and clay). Four to eight grams of wet sediment samples were mixed with 70 mL of saturated cesium chloride solution and centrifuged at 2000 rpm (1000g) for 10 minutes in 80-mL glass centrifuge tubes. The light particles that floated to the top were decanted and collected on a 0.7- μ m glass fiber filter paper. This procedure was repeated 3-5 times until the light fraction was separated from the heavier mineral fraction. Separated fractions were washed with deionized water thoroughly to remove the residual cesium chloride. The mass of each fraction was recorded after air-drying and desiccating for 12 hours. Duplicate tests were performed with each sediment sample with relative standard deviations (RSDs) in all cases less than 2-10 %.

PAH Extraction, Cleanup, and Analysis by GC-FID. The PAH concentrations on bulk sediment samples and separated fractions were measured by following EPA standard method 3550B. Three grams of solid samples were placed in a beaker containing 50 mL hexane/acetone (1:1) mixture and extracted using a Fisher 550 Sonic Dismembrator (Pittsburgh, PA) for 6 minutes (pulsing for 15 s on and 15 s off). The sample was extracted three times and then the extracts were combined, concentrated and changed into solvent cyclohexane. Cleanup was performed on the final extract using an activated silica gel column as outlined in EPA standard method 3630C. An Agilent gas chromatograph 6890N system (Sunnyvale, CA) with a fused silica capillary column (HP-5, 30 m long x 0.25 mm I.D.) and a flamed ionization detector was used for analysis based on EPA standard method 8100 for PAHs. Duplicate extractions were performed for each sediment sample as well as the separated fractions. A blank solvent and a standard solution of 16 EPA PAHs spiked with the internal standard were run with every sequence of samples to assess GC performance and validate calibration.

Organic Carbon (TOC) Measurement. The organic carbon content in each sediment sample was determined by AGVISE Laboratories, Northwood, ND, using the conventional combustion technique followed by nondispersive infrared detection after treating the samples with hydrochloric acid to remove the inorganic carbon that may exist in the solid matrix.

Total Extractable Organics (Oil and Grease). A similar experimental protocol to SW-846 EPA standard method 9071A (Hexane Extractable Material) was employed to quantify the level of oil and grease present in all sediment samples. The only difference is use of an ultrasonic extraction instead of a soxhlet extraction procedure, and a hexane/acetone (1:1, v/v) mixture as a solvent instead of pure hexane. It has been determined that TEO values obtained with this protocol are very close to those from the standard method, with differences less than 5%.

Aqueous Equilibrium Tests. Several different techniques for conducting aqueous equilibrium tests were employed in this project. Except for the non-aqueous phase liquid sample (CR10-NAPL-1523), a polyoxymethylene – solid phase extraction (POM-SPE) method developed by Jonker (6) was adopted to determine the distribution coefficients of PAHs between sediment samples and water. In order to confirm results from POM-SPE experiments, six sediment samples were selected to follow a conventional, direct-contact experimental protocol with alum flocculation to measure the PAH aqueous equilibrium concentrations. This direct contact experimental protocol with flocculation has been performed in several studies in our laboratory (5, 8) and is reliably demonstrated to accurately measure aqueous phase concentrations of PAHs and PCBs from soil and sediment samples without the interference from colloids and incomplete phase separation. For sample CR10-NAPL-1523, however, neither of these two methods works easily because of the presence of an oil emulsion in the aqueous phase. Therefore an air-bridge system was used for this sample, which eliminates any potential oil contamination in the measurement of aqueous equilibrium concentrations.

Prior to experiments, an aqueous phase was prepared with Milli- $Q^{\text{®}}$ water by adding in 100 mg/L sodium azide to inhibit microbial degradation and 0.01 M calcium chloride to compensate the ionic strength in the system. In order to avoid photolysis of PAH compounds, all experiments were carried out in amber glass bottles or covered glassware.

1) Aqueous equilibrium tests with POM-SPE method. Polymethylene (POM) strips were deployed in the sediment-water system as a passive equilibrium sampler. POM strips take up dissolved organic compounds such as PAHs from the sediment-water slurry until phase equilibrium is reached in the system. At equilibrium, the measured PAH concentrations in POM strips, together with the knowledge of concentration-independent POM-water partition

coefficients predetermined for PAH compounds, can be used to quantify the PAH compounds' freely dissolved concentrations.

Prior to aqueous equilibrium experiments with POM-SPE for sediment samples, sorption isotherm tests were conducted in our laboratory to characterize the sorption properties of POM and to determine the POM-water partition coefficients for PAH compounds of interest. Each sorption isotherm comprised seven data points with a span of approximately two orders of magnitude in aqueous concentrations. For more hydrophobic compounds (> benz[a]anthracene) with their extremely low aqueous solubility, the lowest data points of aqueous concentrations were below the limit of aqueous quantification in our study, therefore six pairs rather than seven pairs of aqueous concentrations (C_w) and concentrations sorbed into POM (C_{POM}) are involved in the determination of K_{POM-w} values for these compounds. It was found for all PAH that sorption to POM follows a linear isotherm within aqueous concentration ranges that were examined. The POM-water partition coefficients are defined as the ratio of the compound concentration in the POM (C_{POM}) divided by the aqueous phase concentration (C_w) at equilibrium:

$$K_{POM-w} = C_{POM} \, (\mu g/kg) / C_w \, (\mu g/L) \qquad [L/kg]$$

Due to the linearity of sorption isotherms, the values of K_{POM-w} were determined by taking slopes of the linear regression curves fitted to the sorption isotherm data. Results of sorption isotherm tests confirmed that sorption of PAHs to the POM can be regarded as a partitioning process and is independent of the solute concentration and other organic analytes in the aqueous phase. These properties make this material an excellent passive sampler for quantifying the freely dissolved concentrations of PAH compounds from environmental matrixes. Kinetic studies of PAH uptake into POM strips from water were also performed. Results showed that the attainment of equilibration for PAH uptake by POM requires less than 40 days with more hydrophobic compounds taking less time, and with higher mass ratio of POM-to-water phase taking less time. The detailed results and discussions about the POM-SPE method and applications in environmental analytical measurement were summarized in a recent research paper (7).

The aqueous equilibrium tests with POM strips were set up as follows: 3 g of wet sediment sample, 0.5 g of POM strip, and 35 mL of prepared aqueous phase were placed in a 40 mL glass

bottle with an aluminum-lined lid. Duplicate tests were carried out for each sediment sample. All the bottles were tumbled end-to-end at 4 rpm in the dark at room temperature for a period of 40 days. The mass of POM strip was chosen so that less than 5% of the individual PAH compounds were accumulated into the POM from the water. This ensures that the original equilibrium between sediment and water will not be disturbed by POM as an additional sorbing phase (7, 9).

After equilibration, the POM strips were taken out of the system with a pair of tweezers, rinsed with deionized water and dried with a Kimwipe, and then soxhlet extracted continuously with 150 mL hexane/acetone (1:1, v/v) for 16 hours. An internal standard (2 μ g 2-fluorobiphenyl) and a surrogate PAH compound (1 μ g pyrene- d_{10}) were added prior to extraction. The extracts were then concentrated using a rotoevap followed by a stream of gentle nitrogen purge to 1 mL, and cleaned up for GC-FID analysis as described above. The measured PAH concentrations in POM strips allowed us to calculate the aqueous equilibrium concentrations based on the knowledge of POM-water partition coefficients.

2) Aqueous equilibration tests with batch mixing and particle separation using alum flocculation. In this method, the wet sediment sample and the synthetic aqueous phase in the weight ratio of 1:20 were placed in 1-L glass bottles with a Teflon-lined cap and gently agitated on a roller at 2 rpm in the dark for two weeks to allow complete mixing and full contact between sediment and water. In order to ensure aqueous equilibrium and phase separation within the sediment-water system, these bottles were equilibrated further for two months before sampling. After the sediment fraction settled to the bottom of the glass bottle, 750 mL of the aqueous phase was transferred to a centrifuge tube. To remove particulates from solution without sacrificing the aqueous PAH composition integrity, the tube contents were centrifuged at 20 °C and 1000 g for 10 minutes to settle any coarse particles (5, 8). This step was followed by the addition of about 10 mL 0.1 M alum solution and adjustment of pH back to neutral with 1.0 N NaOH solution. The supernatant water was mixed carefully using a glass pipette for 1 minute to mix and flocculate the alum without disturbing the settled particles. A sweep floc was formed, which co-precipitates with fine particles. The flocculated system was subjected to a second centrifugation step at 500 g for 30 minutes. The clear supernatant created afterwards was

carefully pipetted into a separatory funnel and extracted with fresh volumes of hexane three times. The pipette was similarly rinsed with hexane into the separatory funnel. The hexane extracts were combined, dried using anhydrous sodium sulfate, concentrated to 1 mL, cleaned by a silica gel cleanup method, and the final clean eluate concentrated to 0.2 mL for GC analysis.

Our previous work showed that particulates are effectively precipitated by alum flocs and that aqueous PAH concentrations are not affected by the technique (5). In the absence of particulate removal using alum flocculation, equilibrium tests may be biased because the apparent aqueous phase concentration may comprise both dissolved PAHs and PAHs sorbed on micro-particulates or colloids. This is an especially important concern when measuring PAH concentrations at sub-microgram per liter concentrations as in this study.

3) Aqueous equilibrium tests with an air-bridge system. This test protocol involves an equilibration technique in which an air bridge is employed to physically separate the sediment from the equilibrated water. The air-bridge method described by Bucheli and Gustafsson (10) was used. This method employs a beaker holding the sediment sample that is contained within a larger, closed vessel with an aqueous phase in the outer annular space. PAHs in sediment in the inner beaker are allowed to volatilize and transfer to the outer water phase through the air bridge, and over time an equilibrium state is attained between the sediment slurry inside the beaker and the water in the annular space outside the beaker. For PAHs, the equilibration takes one to two months. In this project, a 150-mL glass beaker containing about 10 grams of sediment sample (CR10-NAPL-1523), 120 mL synthetic aqueous phase and a stir bar was placed in a 4 L glass jar with Teflon-lined cap, and the annular space between the jar and the beaker was filled with 1 L synthetic aqueous phase. The glass jar was sealed with Teflon tape and equilibrated for two months on a magnetic stirrer to gently mix the solid slurry and facilitate solute transfer until sampled. The aqueous phase sampled from the outer annular space was extracted by hexane three times, and then the extracts were combined, concentrated and analyzed by GC for PAHs as described above. The main advantage of the air-bridge method is the absolute certainty of the elimination of particulate-phase contamination in the measurement of aqueous phase concentrations.

Quality Assurance and Quality Control (QA/QC). All solvents used in the experiments

were checked periodically with GC-FID for any possible contamination. Prior to the extraction of sediment samples, a surrogate PAH compound (1 μ g pyrene- d_{10}) and an internal standard (2 μ g 2-fluorobiphenyl) were spiked in the sample matrix to monitor the performance of all procedures involved in PAH analyses in sediment samples. Duplicate tests were carried out with each sample. The reported values are average values with relative standard deviations ranging from 2% to 10%, which are well within the method control limits. A standard solution of 16 EPA PAHs with a known concentration of individual compounds was chosen as a laboratory control sample (LCS), and this was included in each analytical batch and its recovery was used to check the column cleanup efficiency and GC performance. A reagent blank consisting of the elution solvents was passed through the silica gel column and analyzed before each new lot of adsorbents and solvents were used in this method. This was done to ensure that any interferences from adsorbents (silica gel) are well below the method detection limits. The recovery rates of laboratory control samples (LCS) were $93 \pm 7\%$ (n=160) for all PAH compounds of interest, considerably higher than the requirement of $\geq 85\%$ specified in the EPA standard method. The recovery check of the surrogate PAH compound and the internal standard in the column cleanup procedure was also performed, with recovery rates of 95 \pm 8% (n=18) and 96 \pm 4% (n=18) respectively.

A blank reagent and a standard solution spiked with known amount of the internal standard and the surrogate compound were run with each sample batch for GC-FID analysis. The instrumental performance was checked everyday and maintained by following the standard operating procedure (SOP) established in our laboratory, such as regularly changing the inlet liner and the septa for the injection port. GC calibration was performed every three months with standard solutions of 16 EPA PAHs at five different concentration levels. Calibration curves for each PAH compound of interest, which delineate the relationship between instrument responses to concentrations of target compounds in the standard solutions, were constructed and used to calculate the analyte concentrations in samples. GC calibration was verified for each analytical batch based on the instrument responses for the standard solution, making sure to meet the calibration verification criteria of within $\pm 20\%$ difference from the initial calibration responses.

In aqueous equilibrium tests using the POM method, the extraction efficiency of POM strips

with hexane-acetone (1:1, v/v) was found to be $98 \pm 1.4\%$ (n=160) for all 16 EPA PAH compounds. And in the tests using alum-flocculation and air-bridge techniques, PAHs in the aqueous phase were extracted in accordance with EPA standard method 550, and the average recovery rate was determined to be $99 \pm 2\%$.

RESULTS AND DISCUSSIONS

Experimental results for this study are presented and discussed with respect to each of twenty sediment samples from the following three aspects: (a) physical and chemical characteristics, (b) PAH analyses for bulk and separated fractions of sediment samples, and (c) PAH aqueous equilibrium concentration measurements and calculation of site-specific sediment-water distribution coefficients.

Physical and Chemical Characteristics of Sediment Samples. Table 1 summarizes the physical and chemical properties of sediment samples examined in this study, including total organic carbon content, oil and grease content, mass percentage of light density materials, and weight ratio of total organic carbon to light density fraction in the sediment samples.

The total organic carbon (TOC) contents of the sediment samples are highly variable, with the lowest value of 0.7% by weight in sample NLU 402 and the highest value of 77.1% by weight in sample NLU 65. Among these sediment samples, the majority (i.e., thirteen out of twenty) have TOC levels of approximately 14% to 27%. Three samples including NLU 402, NLU 55, and NLU 56 have much lower TOC levels of less than 5% by weight. By contrast, the remaining four of twenty sediment samples have relatively high TOC levels ranging from 40.3% up to 77.1%, which include NLU 51, NLU 65, NLU 73-Stanford, and the NAPL sample, CR10-NAPL. The mass of light fraction particles (density < 1.8 g/cc) determined by density separation varied from sample to sample, ranging from less than 5% to 79% by weight.

The oil and grease contents of sediment sample varied from 0.2% to 2.5%, except the sample CR10-NAPL, which appears to be very oily, soft-tar like, having an oil and grease content of greater than 95% by weight. Oil and grease is a conventional pollutant defined under 40 CFR 401.16 by US EPA and generally refers to substances, including biological lipids and mineral

hydrocarbons that have similar physical characteristics and common solubility in an organic extracting solvent. By comparing the oil and grease content with the PAH analysis results described below, we note that sediment samples with high TOC content have relatively higher oil and grease content and correspondingly higher levels of PAH contamination. The examples are the four samples with relatively high TOC values, NLU 51, NLU 65, NLU 73-Stanford, and CR10-NAPL.

Organic particles such as plantative materials and black carbons are generally less dense than the mineral (sand/clay/silt) fractions of sediments. Thus, low density fractions obtained by the density separation technique from sediment samples tend to comprise the majority of organic particles. Given the organic carbon content in a certain class of organic particles, the mass ratio of total organic carbon to the less-dense particulate fraction could provide a quick assay to predict the composition of organic carbon in the sediment sample. Our experience is that if the light density fraction comprises primarily black carbonaceous materials, then owing to their extremely high organic carbon content, usually close to or over 90% by weight, the TOC values and the light fraction measurements are in the same range for a ratio of TOC to light fraction in the proximity of unity. This means that the organic carbon in these sediments is mainly contributed by black carbon matter, and examples from this study are NLU 51, NLU 65 and NLU 73-Stanford. On the other hand, for cases as NLU 44, NLU 47 and NLU 62 etc., with TOC values around 15 ~ 17%, the light (density < 1.8 g/cc) fraction accounts for about 39 ~ 46% of total sample weight. This indicates that the weight percent of carbon in the less-dense particulate fraction for these samples is about 40%, which is expected if the majority of the organic particulate matter is plantative material. We also note that three samples NLU 68-S1, NLU 68-S2 and NLU 68-SS have a TOC-to-light fraction ratio of 0.6 or so, and based on the same reasoning, this suggests that the light fraction could be a mixture of both black carbonaceous materials and natural organic matter with either of these two components dominating.

PAH Analysis and Distribution Patterns. The experimental results of PAH analyses in bulk sediment samples, as well as that for density separated fractions, are summarized in Table 2 and Table 3. The detailed data for individual PAH concentrations and distributions among the

light density fraction and the heavy density fraction for each sample are presented in Appendix I, except for the NAPL sediment sample for which the density separation is not applicable. As shown in Table 2, all twenty sediment samples exhibit various degrees of PAH contamination and the total 16 EPA PAH concentrations range from 18 mg/kg for NLU 54 to 4,390 mg/kg for NLU 51. The total 16 EPA-listed PAHs content of the NAPL sample is 90,480 mg/kg. Figures 1 and 2 depicted the distribution profiles of the 16 EPA PAHs in six highly or moderately contaminated sediment samples. Profiles for NLU 51, NLU 65, and NLU 73-Stanford are shown in Figure 1 for which total PAH concentrations range from 1,900 to 4,400 mg/kg, and profiles for NLU 56, NLU 64, and NLU 68-SS are shown in Figure 2 with total PAH levels in the range of 850 to 1,400 mg/kg. As shown in the figures, despite the various PAH concentration levels, a common feature for these samples is that fluoranthene and pyrene are the most abundant PAH compounds, except NLU 73-Stanford which has naphthalene and phenanthrene as dominant PAHs.

PAH concentrations on lighter and heavier fractions of each sediment sample were measured to examine the association and distribution patterns within the sample. The total concentrations of 16 EPA-listed PAHs on light-density and heavy-density fractions as well as the mass percent of PAHs associated with the less-dense fraction are reported in Table 3. From the data summarized in Table 3 and Appendix I, it is evident that PAH concentrations in the lighter density particle fractions are over four times greater than those in the heavier density particle fractions. In the case of NLU 65, the total concentration of PAHs in the light density particle fraction is 194-fold greater than the heavier fraction. Based on the knowledge of PAH concentrations in bulk samples, as well as separated fractions and measurements of mass percentage of light fraction for each sediment sample, it can be estimated that for most of these sediment samples the majority of the PAHs (78 ~ 100% of total mass) are associated with the low-density particle fraction. Only two samples, NLU 402 and NLU 68-US-S2, have lower values of 69% and 61% respectively for the mass of PAHs in the low-density fraction. The total PAH concentration in the light density particle fraction of sample NLU 402 is 2,905 mg/kg, which is 45 times as high as the concentration value (64 mg/kg) in the heavy density fraction. Since NLU 402 is the sample with the lowest TOC value (0.7% by weight) and the lowest light fraction (4.6% by weight), it is not surprising to observe that relatively less mass of PAHs are in the light density fraction.

Sediment-water Distribution Coefficients (K_d) for PAHs. Using the knowledge of the equilibrium partitioning of PAHs between POM strips and water established earlier (7), the aqueous equilibrium concentrations of PAHs from sediment samples were calculated based on measurements of PAH concentrations in POM strips according to $C_w = C_{POM} / K_{POM-water}$. From this, sediment-water distribution coefficients K_d were derived using the measured sediment PAH concentrations and the dissolved PAH concentrations inferred from the POM samplers. $K_{POM-water}$ values for 12 PAH compounds determined in our previous studies are listed in Table 4, along with literature values of octanol-water (K_{ow}) partition coefficients and organic carbon-normalized partition coefficients (K_{oc}). Values of K_{ow} were taken from the reference text *Environmental Organic Chemistry* by Schwarzenbach et al. (15), except that K_{ow} values of benzo[b]fluoranthene, benzo[k]fluoranthene and dibenz[a,h]anthracene are taken from Accardi-Dey et al. (16), and the K_{ow} value of benzo[g,h,i]perylene is taken from Mackay et al. (12). K_{oc} values are calculated from the equation:

$$\log K_{oc} = 0.989 \log K_{ow} - 0.346$$

 $K_d = f_{oc} K_{oc}$ [L/kg]

which was originally developed by Karickhoff et al. (1). This equation is very close to the correlation given in the reference text *Environmental Organic Chemistry* (15): $\log K_{oc} = 0.98 \log K_{ow} - 0.32$ for PAH compounds with $\log K_{ow}$ values of 2.2 to 6.4. In the nineteen consecutive tables in Appendix II, the aqueous equilibrium concentrations determined by the POM-SPE method and the calculated site-specific sediment-water distribution coefficients for individual PAHs are presented for each sample.

In order to verify aqueous equilibrium concentration measurements and K_d values determined by the POM-SPE method, an alum-flocculation technique was applied to six selected sediment samples. Experimental results confirmed that PAH aqueous equilibrium concentrations measured by these two methods as well as K_d values calculated correspondingly agree with each other very well. Figure 3 shows the correlation between logarithmic K_d values obtained with POM-SPE method and alum-flocculation method for samples NLU 73-stanford and NLU 51. Data shown in figure 3 demonstrate that the POM-SPE method can serve as a good alternative to the alum-flocculation method.

We compared our experimentally measured K_d values with the conventional organic-carbon equilibrium partitioning model, in which the K_d value was predicted on the basis of each sediment's organic carbon content (f_{oc}) and each PAH compound's K_{oc} value according to Karickhoff et al. (1) as cited above. This comparison shows that these sediment samples exhibit distinctive partitioning behaviors. Accordingly, three conceptual sorption models are proposed for adoption in the current study as follows:

1. Modified fraction organic carbon partitioning: in this model, all the organic matter present in the sample is assumed to be uniformly distributed and homogeneous in terms of sorption properties, and PAHs sorbed on sediment solids can be regarded as partitioning into the organic carbon domain. With this approach, the organic carbon content (f_{oc}) is considered to be the crucial parameter in the prediction of sediment-water distribution coefficients. In Karickhoff's model, the f_{oc} is considered to be comprised of natural plant and animal derived organic carbon. In the modified model, we envision the f_{oc} to be comprised partly of natural organic carbon and partly of black carbonaceous materials.

Fifteen sediment samples agreed reasonably well with this model, and they include: NLU 44, NLU 47, NLU 402, NLU 54, NLU 55, NLU 57, NLU 62, NLU 64, NLU 68-S1, NLU 68-S2, NLU 72, NLU 73-SS, NLU 45, NLU 58, and NLU 68-SS. These sediment samples have relatively low organic carbon contents (0.7 ~ 26.5% by weight) and low PAH concentration levels (total concentration 18 ~ 320 mg/kg) except NLU 64 and NLU 68-SS with relatively high 16 EPA PAH concentrations of 850 mg/kg and 1120 mg/kg, respectively.

Figure 4 shows a comparison between experimentally measured PAH sediment-water distribution coefficients and model predictions for two representative samples (a) NLU 44 and (b) NLU 64. As demonstrated in the diagrams, a common trend we find from the comparison between observed K_d values and model-predicted K_d values is that the observed site-specific K_d values are generally larger than those predicted by fraction organic carbon (OC) partitioning by

approximately $0.5 \sim 1.0$ logarithmic units, with the divergence from the model greater for the more hydrophobic PAH compounds. This implies that other than natural organic matter, some other types of organic carbon particles, such as black carbonaceous materials, affect the sorption process due to their higher sorption capacity and non-linear equilibrium sorption mechanisms for PAHs. The presence of black carbonaceous particles in these sediment samples is expected since the sediment samples examined in this study originated from the lake area surrounding the former manufactured gas and tar refining plant sites, and received runoff that likely contained soot from the surrounding urban areas.

2. Oil/tar partitioning: this model assumes that all PAHs are present in an oil phase and that the partitioning behavior follows dissolution from the oil phase. Our previous study (5) used an oil/tar-water partitioning model to describe PAH partitioning from lampblack-impacted soil samples governed by an oil/tar phase. In this approach the aqueous equilibrium concentration of a PAH compound is calculated using the equation:

$$C_{aq,i} = x_i S_i$$

where x_i is the mole fraction of PAH *i* in the oil/tar phase, and S_i is the subcooled liquid solubility of PAH *i*. The subcooled liquid solubility of PAH *i* is calculated by dividing the aqueous solubility of the solid PAH *i* by the ratio of the solid-liquid reference fugacities $(f^{s}/f^{t})_{i}$ for the pure PAH *i*, as summarized by Peters et al. (13). The mole fraction of PAH *i* is computed by assuming that the oil/tar phase is measured as oil and grease (total extractable organics) with an average molecular weight of 270.

In the current study, this model was applied to three sediment samples (NLU 51, NLU 65 and NLU 56) and the NAPL sample to interpret PAH sorption mechanisms and partitioning properties. Figure 5 shows a comparison of the apparent PAH sediment-water distribution coefficients determined in this study with different model predictions for samples (a) NLU 51 and (b) NLU 65. It was noted that the fraction organic carbon model and oil/tar model are comparable in predicting the sediment-water distribution coefficients of PAHs for these two samples. This is because that the weight ratios of oil and grease content to TOC are 0.06 and 0.03 for NLU 51 and NLU 65 respectively, and the corresponding sorption capacity of oil and grease for PAHs is

generally greater than that of natural organic matter by a factor of 5 to 25. Thus, these two factors approximately compensate each other and the predicted PAH distribution coefficients on the basis of these two models result in very similar values. Nonetheless, we believe that the oil/tar model is more appropriate for the following reasons. These two sediment samples have very high TOC values of 40.3% for NLU 51 and of 77.1% for NLU 65. The fact that the ratio of TOC value to light-density fraction is close to or greater than unity implied that black carbon materials may be the dominant type of organic carbon in these samples. NLU 51 and NLU 65 are two highly contaminated sediment samples with the highest total PAH concentration levels at 4,400 mg/kg and 4,200 mg/kg respectively, which may exhaust the black carbon sorption capacity. Based on these characteristics, it is conceivable that oil tar-water partitioning model would be more appropriate to interpret PAH sorption behavior in NLU 51 and NLU 65 than a fraction It was observed that aqueous equilibrium organic carbon-water partitioning model. concentrations measured with these two samples are significantly higher than that for the sediment sample controlled by soot carbon. For example, freely dissolved aqueous concentration of phenanthrene from NLU 51 is 103 μ g/L and 22 μ g/L for NLU 65, as opposed to 0.5 µg/L for NLU 73-Stanford which is controlled by oil soot partitioning as explained below.

As shown in figure 6, PAH partitioning behavior in sediment sample NLU 56 is described by the oil/tar-water partitioning model. NLU 56 has a very low TOC value of 2.7% but a relatively high PAH concentration at 1360 mg/kg. These factors result in the oil/tar-water partitioning model providing a much better fit to experimental results, especially for more hydrophobic PAH compounds.

Due to its unique nature as a soft-tar like material, the PAH aqueous equilibrium concentrations from sample CR10-NAPL can be predicted very well using the oil tar-water partition model. Figure 7 shows the good correlation between the measured distribution coefficients with the model predictions.

3. Soot-carbon controlling partition model: in this model, black carbon is the predominant form of organic particulate matter in the sediment, and PAHs are assumed to be associated with the soot carbon. Soot carbon normalized distribution coefficient K_{oc} values of PAHs were determined in our previous work with lampblack-impacted soils. As presented in

Table 4, these values are generally $1.5 \sim 2.5$ orders of magnitude larger than those derived from natural organic matter due to the strong affinity of soot carbon for PAHs. Therefore a striking phenomenon often observed with this kind of sample is the diminished aqueous equilibrium concentrations and drastically elevated sediment-water distribution coefficients (K_d). NLU 73-Stanford is the only sample among this batch of sediments that clearly exhibits this unique partitioning behavior for PAHs.

Figure 8 shows that the aqueous equilibrium concentrations of PAHs from NLU 73-Stanford are in good agreement with the predictions of a soot partitioning model. NLU 73-Stanford has a total organic carbon content of 74.8% by weight, and this is very close to the light density fraction of 72.9% determined by density separation. The fact that the weight ratio of TOC versus light fraction is nearly unity suggests that black carbon is the major organic carbon source in this sample.

Based on the experimental results and modeling approaches demonstrated above, it is noted that even though sediment samples have similar compositions of organic matter with black carbonaceous particles being present, these samples may exhibit two different PAH partitioning behaviors depending on the predominance of either oil/tar or soot-carbon partitioning. We postulate that these phenomena can be explained on the basis of physiochemical characteristics of the black carbon solid matrix and sorptive nature of PAHs. The strong binding of PAHs to a black carbon solid matrix is attributed to the overlap of π -electrons in the aromatic soot structure and the planar aromatic rings of PAH molecules. Soot carbon has a non-linear equilibrium sorption isotherm, and the amount of PAHs that can be sorbed onto the soot carbon skeleton is limited. When the PAH concentrations reach the sorption capacity of soot carbon, they start to form a free oil phase, and the sorption mechanism for PAHs undergoes a transition from adsorption on a soot carbon matrix to absorption into a free aromatic oil phase, therefore the aqueous partitioning behaviors for PAHs in these kind of samples appear like dissolving from oil tar phase rather than from the strong carbon sorbent.

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Sample ID	TOC (%)	Carbonate (%)	Oil & grease (%)	Light fraction (%)	TOC vs. light fraction
NLU 44	14.8		0.98	38.7 ± 3.8	0.4
NLU 47	14.1		1.25	45.6 ± 1.3	0.3
NLU 51	40.3		2.44	44.9 ± 3.3	0.9
NLU 402	0.7		0.17	4.6 ± 0.7	0.2
NLU 54	15.2		0.92	45.0 ± 1.7	0.3
NLU 55	3.8		0.15	13.8 ± 0.5	0.3
NLU 56	2.7	0.2	0.60	9.2 ± 0.8	0.3
NLU 57	14.1	0.1	0.97	47.0 ± 2.2	0.3
NLU 62	16.8		1.27	46.0 ± 1.9	0.4
NLU 64	18.2	0.1	1.20	46.0 ± 0.7	0.4
NLU 65	77.1	0.1	2.15	46.4 ± 1.3	1.7
NLU 68-S1	15.8	0.1	1.38	25.7 ± 2.5	0.6
NLU 68-S2	14.6	0.2	1.06	23.0 ± 2.1	0.6
NLU 72	16.6		0.86	47.0 ± 3.0	0.4
NLU 73-SS	15.4		1.22	46.2 ± 1.8	0.3
NLU 73-Stanf	74.8		1.55	72.9 ± 1.6	1.0
NLU 45	13.5	0.1	0.98	56.1 ± 0.5	0.2
NLU 58	16.2	0.1	1.44	78.8 ± 3.7	0.2
NLU 68-SS	26.5		2.49	43.8 ± 0.7	0.6
CR10-NAPL	56.2		> 95%	n.a.	n.a.

Table 1 Characteristics of Lake Union sediment samples, including total organic carbon(TOC), oil and grease content, and weight percentage of light density fraction

Compounds	Sample ID										
_	NLU 44	NLU 47	NLU 51	NLU 402	NLU 54	NLU 55	NLU 56	NLU 57	NLU 62	NLU 64	NLU 65
Naphthalene	0.2	0.7	19	2.3	n.d.	4.2	3.2	n.d.	3.1	3.0	14
Acenaphthylene	n.d.	0.4	16	4.6	n.d.	1.7	6.3	n.d.	1.4	4.2	27
Acenaphthene	n.d.	0.5	69	14	n.d.	1.8	37	0.2	3.3	12	64
Fluorene	n.d.	0.5	44	13	0.2	1.8	15	1.0	2.4	6.2	27
Phenanthrene	1.2	3.9	501	59	1.1	12	131	3.0	11	31	174
Anthracene	0.3	0.8	160	17	0.3	2.6	35	2.1	3.3	16	48
Fluoranthene	3.5	11.1	851	28	2.4	21	240	8.3	22	136	690
Pyrene	4.7	13.7	1009	32	2.8	25	300	7.0	28	169	908
Benz[a]anthracene	1.6	4.1	236	11	1.0	8.4	65	2.8	8.4	47	242
Chrysene	1.7	3.9	271	11	1.1	9.7	79	4.0	9.2	52	291
Benzo[b]fluoranthene	2.5	4.9	202	5.1	1.5	7.5	66	3.2	9.6	56	279
Benzo[k]fluoranthene	3.3	5.8	198	6.0	2.8	7.9	66	4.6	11	55	270
Benzo[a]pyrene	3.2	7.1	342	9.1	1.2	11	114	3.2	15	93	439
Indeno[1,2,3-cd]pyrene	2.5	5.4	205	4.1	1.6	7.6	90	3.1	13	76	333
Dibenz[a,h]anthracene	0.2	0.6	26	0.9	n.d.	1.2	11	0.4	1.7	8.9	47
Benzo[g,h,i]perylene	2.7	5.7	235	3.8	1.5	8.2	104	3.2	15	90	382
Total 16 EPA PAHs	28	69	4390	220	18	131	1360	46	157	854	4235

 Table 2 Summary of PAH concentrations in Lake Union sediment samples (mg/kg dry weight sediments)

Compounds	Sample ID								
_	NLU 68-S1	NLU 68-S2	NLU 72	NLU 73-SS	NLU 73-Stanford	NLU 45	NLU 58	NLU 68-SS	CR10-NAPL
Naphthalene	28	28	0.7	5.0	326	1.0	3.2	3.8	22890
Acenaphthylene	2.4	4.2	0.6	3.5	17	1.2	3.2	5.4	613
Acenaphthene	6.6	3.9	0.6	6.3	150	3.0	2.2	32	6700
Fluorene	4.2	7.1	0.3	3.9	71	1.4	2.2	21	2450
Phenanthrene	24	26	3.2	26	362	9.8	20	45	14380
Anthracene	6.6	6.0	0.8	7.6	86	2.6	4.6	43	2769
Fluoranthene	42	13	14	45	219	38	30	193	8580
Pyrene	55	19	17	57	276	46	37	244	11100
Benz[a]anthracene	19	6.7	5.6	18	60	14	13	65	2834
Chrysene	23	8.4	5.8	23	75	15	16	77	2830
Benzo[b]fluoranthene	19	3.5	7.7	19	48	16	13	59	2287
Benzo[k]fluoranthene	11	4.4	8.5	22	52	17	15	63	2560
Benzo[a]pyrene	27	5.4	12	31	74	26	19	101	4230
Indeno[1,2,3-cd]pyrene	22	3.2	11	24	50	21	16	76	2792
Dibenz[a,h]anthracene	3.5	0.9	1.2	3.4	12	2.6	2.5	11	430
Benzo[g,h,i]perylene	25	3.4	12	26	55	23	18	86	3030
Total 16 EPA PAHs	316	142	100	319	1932	237	214	1124	90480

 Table 2 (continued)
 Summary of PAH concentrations in Lake Union sediment samples (mg/kg dry weight sediments)

Sample ID	Total 16 PAHs concentration in light density fraction (mg/kg dry weight)	Total 16 PAHs concentration in heavy density fraction (mg/kg dry weight)	Mass percentage of PAH in light density fraction (% by weight)
NLU 44-SS-0010	47	8	78
NLU 47-SS-0010	118	10	91
NLU 51-SS-0010	8738	542	93
NLU 402-GE15	2905	64	69
NLU 54-SS-0010	28	2	93
NLU 55-SS-0010	868	8	95
NLU 56-SS-0010	12779	181	88
NLU 57-SS-0010	63	13	81
NLU 62-SS-0010	296	29	90
NLU 64-SS-0010	1697	128	92
NLU 65-SS-0010	8937	46	99
NLU 68-US-S1	1119	67	85
NLU 68-US-S2	334	63	61
NLU 72-SS-0010	165	28	84
NLU 73-SS-0010	800	11	98
NLU 73-Stanford	2561	183	97
NLU 45-DC	457	49	92
NLU 58-SS-0010	340	12	99
NLU 68-SS-0010	2584	259	89
CR10-NAPL	n.a.	n.a.	n.a.

Table 3 Summary of PAH analyses on Separated Fractions

Note:

* Please refer to Appendix I for detailed data of PAH distribution on separated fractions for each sample.

Table 4

			1	1
Compounds	log K _{ow} ^[1] (-)	log K _{oc} ^[2] (L/kg)	log K _{oc-lampblack} ^[3] (L/kg)	log K _{POM-w} ^[4] (L/kg)
Naphthalene	3.33	2.95		2.06
Acenaphthylene	4.20	3.81		n.a.
Acenaphthene	4.00	3.61		2.43
Fluorene	4.32	3.93		2.88
Phenanthrene	4.57	4.17	5.57	3.22
Anthracene	4.68	4.28	5.91	3.46
Fluoranthene	5.23	4.83	6.74	3.73
Pyrene	5.13	4.73	6.81	3.74
Benz[a]anthracene	5.91	5.50	7.34	4.36
Chrysene	5.81	5.40	7.44	4.32
Benzo[b]fluoranthene	5.90	5.49	8.05	4.75
Benzo[k]fluoranthene	6.00	5.59	8.08	4.63
Benzo[a]pyrene	6.13	5.72	8.35	4.73
Indeno[1,2,3-cd]pyrene				n.a.
Dibenz[a,h]anthracene	6.80	6.38		n.a.
Benzo[g,h,i]perylene	7.10	6.68		n.a.

Partition Coefficients Used to Interpret Aqueous Equilibrium Experiments: Octanol-Water (K_{ow}), Organic Carbon-Water (K_{oc}), Lampblack (oil soot) Organic Carbon-Water (K_{oc-lampblack}), and Polyoxymethylene (POM)-Water (K_{POM-w})

Note:

- 1. Log K_{ow} values are taken from the reference *Environmental Organic Chemistry* by Schwarzenbach, Rene P.; Gschwend, P.M. and Imboden, Dieter M.; published by John Wiley & Sons, Inc., Hoboken, NJ, 2003, 2nd edition. Except that log K_{ow} values of benzo[b]fluoranthene, benzo[k]fluoranthene and dibenz[a,h]anthracene are taken from Accardi-Dey, A.; Gschwend, P.M., 2003 (16). Log K_{ow} value of benzo[g,h,i]perylene is taken from Mackay et al. (12).
- 2. Log K_{oc} values are calculated from the equation log $K_{oc} = 0.989 \log K_{ow} 0.346$ established by Karickhoff, S.W. (1)
- Log K_{oc-lampblack} values are determined in a previous study with lampblack-impacted soil samples (5).
- 4. Log K_{POM-water} values are determined in our laboratory for 12 PAH compounds based on extensive sorption isotherm studies with POM strips; POM-water partition coefficients for acenaphthylene, indeno[1,2,3-cd]pyrene, dibenz[a,h]anthracene and benzo[g,h,i]perylene are not currently available (7).

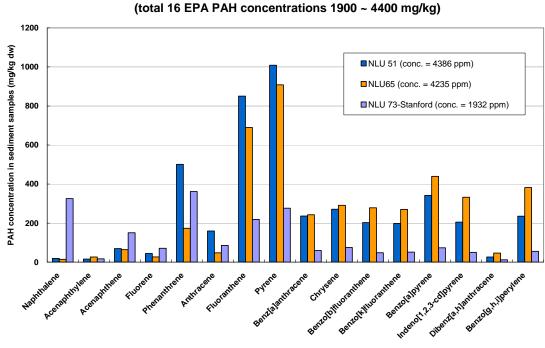
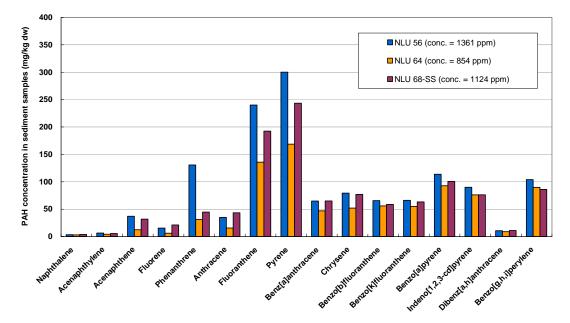


Figure 1 PAH distribution profiles for sediment samples NLU 51, NLU 65 and NLU 73-Stanford (total 16 FPA PAH concentrations 1900 ~ 4400 mg/kg)

Figure 2 PAH distribution profiles for sediment samples NLU 56, NLU64 and NLU 68-SS (total 16 EPA PAH concentrations 850 ~ 1400 mg/kg)



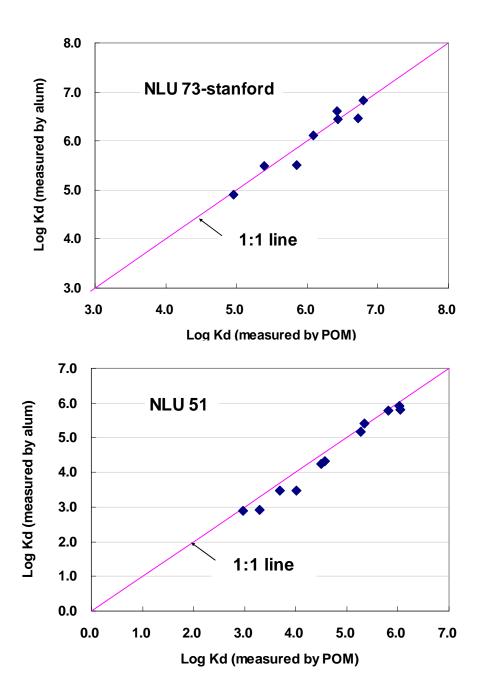


Figure 3 Comparison of sediment-water distribution coefficients (K_d) measured by two different methods, POM-SPE and alum-flocculation, for sample NLU 73-stanford and NLU 51. Experimental results show an excellent agreement in PAH aqueous equilibrium concentrations measured by these two methods, as well as the K_d values calculated correspondingly.

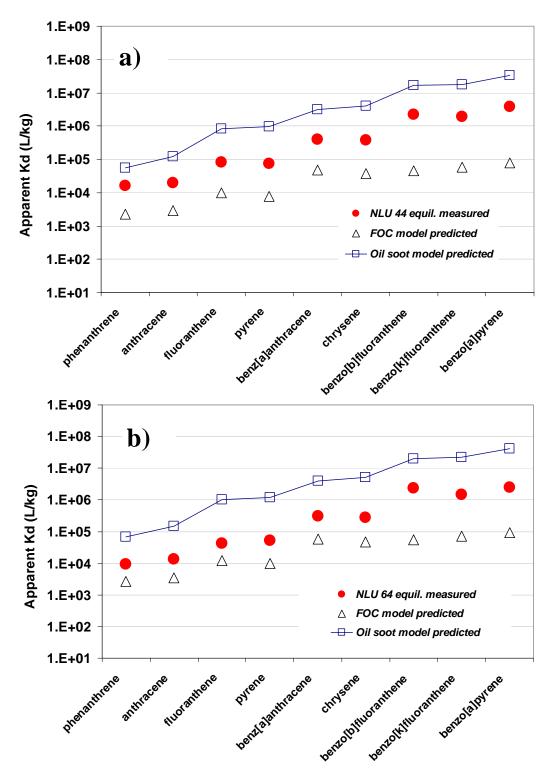


Figure 4 Comparisons between experimentally measured PAH sediment-water distribution coefficients with model predictions (fraction organic carbon partitioning model and oil-soot partitioning model) for two representative samples (a) NLU 44 and (b) NLU 64 in which natural organic matter is considered the dominant type of organic carbon. This representation is characteristic of fifteen sediment samples.

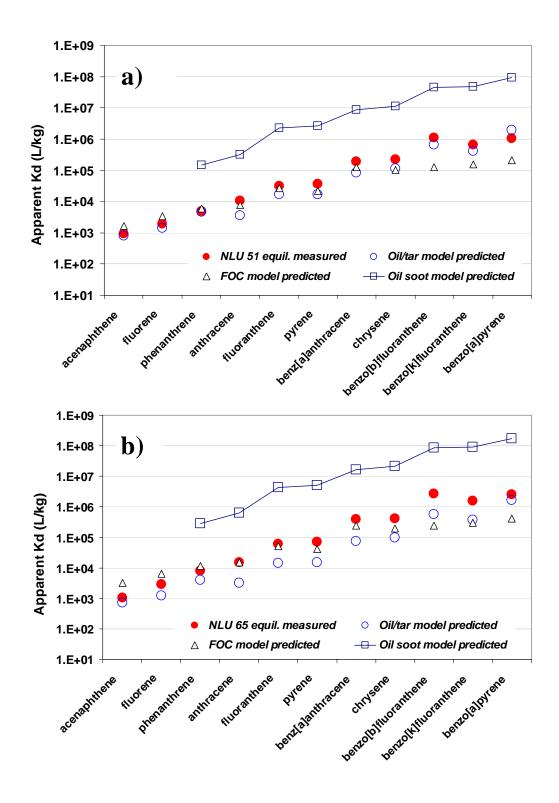


Figure 5 Comparisons between experimentally measured PAH sediment-water distribution coefficients with model predictions for samples (a) NLU 51 and (b) NLU 65 in which the majority of the PAH partitioning is described by equilibrium with an oil/tar phase.

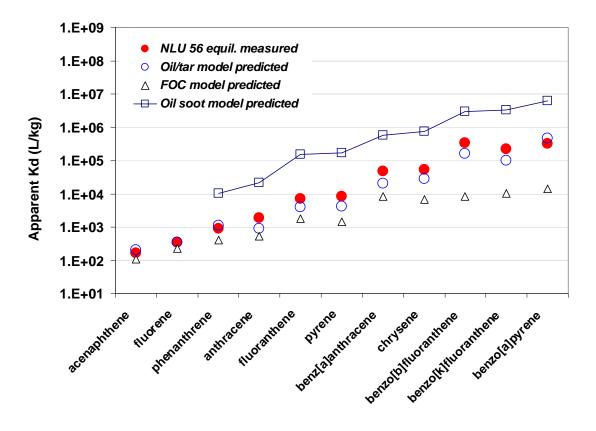


Figure 6 Comparisons between experimentally measured PAH sediment-water distribution coefficients with model predictions for sample NLU 56, showing that the oil tar-water partitioning model better describes the PAH partitioning behavior from this sample than the other models.

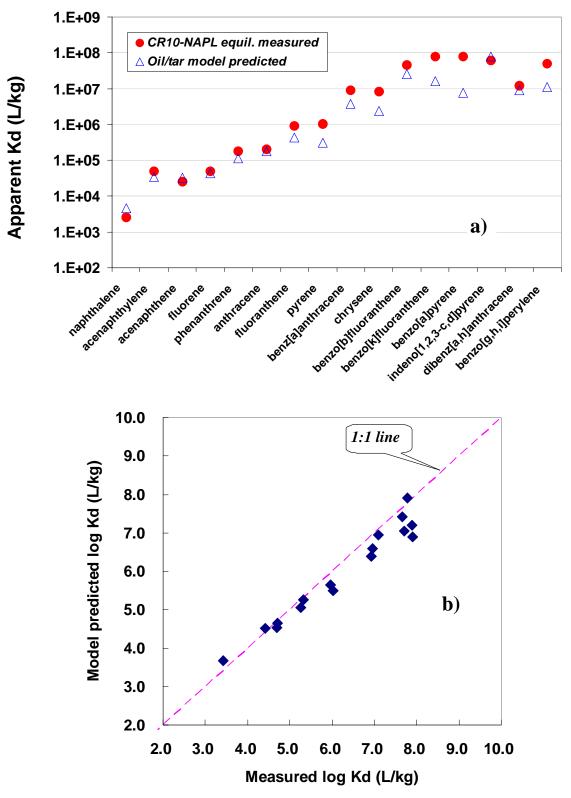


Figure 7 (a) Comparison between experimentally measured PAH sediment-water distribution coefficients with oil tar-water partition model predictions for sample CR10-NAPL. (b) Experimental results are in good agreement with the oil tar-water partitioning model.

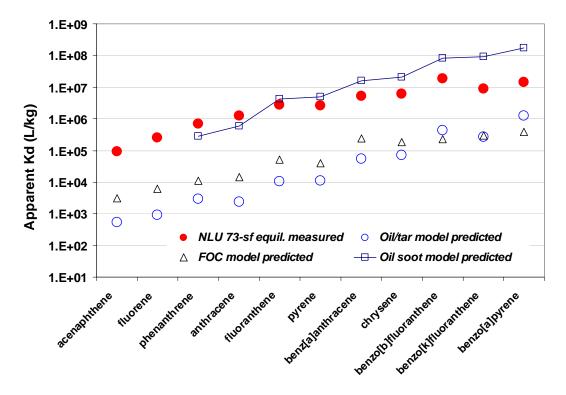


Figure 8 Comparisons between experimentally measured PAH sediment-water distribution coefficients with model predictions for sample NLU 73-Stanford, showing that the oil soot model best describes the PAH partitioning behavior from this sample.

SUB-ATTACHMENT 2D-7.1

Summary of PAH Analyses in Bulk Sediment Samples and Separated Fractions

Compounds	INVERTING ADD. ADD. DAVE DAVE DAVE	Sediment concentration in heavy density fraction (mg/kg dry weight)	Calculated sediment concentration (mg/kg dry weight)	Measured bulk sediment concentration (mg/kg dry weight)	Mass percentage of PAH in light density fraction (% by weight)
Naphthalene	< 0.1	< 0.1	n.a.	0.2	n.a.
Acenaphthylene	< 0.1	< 0.1	n.a.	n.d.	n.a.
Acenaphthene	< 0.1	< 0.1	n.a.	n.d.	n.a.
Fluorene	< 0.1	< 0.1	n.a.	n.d.	n.a.
Phenanthrene	2.2	0.6	1.2	1.2	69
Anthracene	< 0.1	< 0.1	n.a.	0.3	n.a.
Fluoranthene	6.9	1.0	3.3	3.5	82
Pyrene	9.2	1.5	4.4	4.7	80
Benz[a]anthracene	3.1	0.5	1.5	1.6	80
Chrysene	3.2	0.5	1.5	1.7	80
Benzo[b] fluoranthene	3.5	0.8	1.8	2.5	74
Benzo[k]fluoranthene	4.6	0.8	2.3	3.3	77
Benzo[a]pyrene	5.3	0.8	2.5	3.2	81
Indeno[1,2,3-cd]pyrene	4.0	0.9	2.1	2.5	73
Dibenz[a,h]anthracene	< 0.1	< 0.1	n.a.	0.2	n.a.
Benzo[g,h,i]perylene	4.8	0.9	2.4	2.7	77
Total (mg/kg)	46.8	8.3	23.2	27.7	78

NLU 44-SS-0010: Summary of PAH Analyses on Bulk Sediment Sample and Separated Fractions

Note:

n.a. = not applicable, n.d. = not detected or below detection limit, the method detection limit (MDL) for PAH analysis on sediment samples has been determined as 0.1~0.5 mg/kg dry weight for individual PAH compounds.

Reported concentration values are average values of duplicate tests with relative standard deviation (RSD) ranging from 1~17% for all PAH compounds of interest.

Concentration values reported in each fraction are all based on the dry weight of corresponding fractions, for example, concentrations in the light fraction have a unit of mg PAH compounds / kg dry weight of light fraction of this sediment sample.

Compounds	DAVE ELEMENT AND DAVE DAVE DAVE DAVE	Sediment concentration in heavy density fraction (mg/kg dry weight)	Calculated sediment concentration (mg/kg dry weight)	Measured bulk sediment concentration (mg/kg dry weight)	Mass percentage of PAH in light density fraction (% by weight)
Naphthalene	< 0.5	< 0.5	n.a.	0.7	n.a.
Acenaphthylene	< 0.5	< 0.5	n.a.	0.4	n.a.
Acenaphthene	< 0.5	< 0.5	n.a.	0.5	n.a.
Fluorene	< 0.5	< 0.5	n.a.	0.4	n.a.
Phenanthrene	6.4	0.7	3.3	3.9	88
Anthracene	2.0	< 0.5	0.9	0.8	100
Fluoranthene	18.8	1.4	9.3	11.1	92
Pyrene	23.6	1.9	11.8	13.7	91
Benz[a]anthracene	7.4	0.6	3.7	4.1	91
Chrysene	6.7	0.6	3.4	3.9	91
Benzo[b]fluoranthene	7.9	1.1	4.2	4.9	86
Benzo[k]fluoranthene	9.7	0.8	4.9	5.8	91
Benzo[a]pyrene	14.5	0.9	7.1	7.1	93
Indeno[1,2,3-cd]pyrene	10.2	1.1	5.2	5.4	88
Dibenz[a,h]anthracene	< 0.5	< 0.5	< 0.5	0.6	n.a.
Benzo[g,h,i]perylene	11.0	0.9	5.5	5.7	91
Total (mg/kg)	118.1	10.0	59.3	69.2	91

NLU 47-SS-0010: Summary of PAH Analyses on Bulk Sediment Sample and Separated Fractions

Note:

Reported concentration values are average values of duplicate tests with relative standard deviation (RSD) ranging from 0.3~12% for all PAH compounds of interest.

Compounds	Table Contract Least of Contract States South	Sediment concentration in heavy density fraction (mg/kg dry weight)	Calculated sediment concentration (mg/kg dry weight)	Measured bulk sediment concentration (mg/kg dry weight)	Mass percentage of PAH in light density fraction (% by weight)
Naphthalene	40	0.6	18	19	98
Acenaphthylene	32	0.9	15	16	97
Acenaphthene	85	4	41	69	95
Fluorene	65	4	31	44	94
Phenanthrene	922	73	454	501	91
Anthracene	262	16	127	160	93
Fluoranthene	1678	124	822	851	92
Pyrene	2037	149	997	1009	92
Benz[a]anthracene	505	29	243	236	93
Chrysene	553	40	271	271	92
Benzo[b]fluoranthene	476	17	223	202	96
Benzo[k]fluoranthene	396	23	191	198	93
Benzo[a]pyrene	653	27	308	342	95
Indeno[1,2,3-cd]pyrene	447	16	209	205	96
Dibenz[a,h]anthracene	69	2	32	26	97
Benzo[g,h,i]perylene	516	17	241	235	96
Total (mg/kg)	8738	542	4222	4386	93

NLU 51-SS-0010: Summary of PAH Analyses on Bulk Sediment Sample and Separated Fractions

Note:

Reported concentration values are average values of duplicate tests with relative standard deviation (RSD) ranging from 2~24% for all PAH compounds of interest.

Compounds	and matter test of the set and build	Sediment concentration in heavy density fraction (mg/kg dry weight)	Calculated sediment concentration (mg/kg dry weight)	Measured bulk sediment concentration (mg/kg dry weight)	Mass percentage of PAH in light density fraction (% by weight)
Naphthalene	10	0.2	1	2	70
Acenaphthylene	61	1.5	4	5	66
Acenaphthene	114	4.5	10	14	55
Fluorene	112	4.0	9	13	58
Phenanthrene	662	18	48	59	64
Anthracene	211	4.7	14	17	68
Fluoranthene	381	7.5	25	28	71
Pyrene	441	8.4	28	32	72
Benz[a]anthracene	166	3.0	11	11	73
Chrysene	170	3.0	11	11	73
Benzo[b]fluoranthene	86	1.4	5	5	75
Benzo[k]fluoranthene	113	1.9	7	6	74
Benzo[a]pyrene	178	2.9	11	9	75
Indeno[1,2,3-cd]pyrene	89	1.4	5	4	75
Dibenz[a,h]anthracene	24	0.3	1	1	80
Benzo[g,h,i]perylene	87	1.3	5	4	76
Total (mg/kg)	2905	64	195	220	69

NLU 402-GE15-16.5: Summary of PAH Analyses on Bulk Sediment Sample and Separated Fractions

Note:

Reported concentration values are average values of duplicate tests with relative standard deviation (RSD) ranging from 1~19% for all PAH compounds of interest.

Compounds		Sediment concentration in heavy density fraction (mg/kg dry weight)	Calculated sediment concentration (mg/kg dry weight)	Measured bulk sediment concentration (mg/kg dry weight)	Mass percentage of PAH in light density fraction (% by weight)
Naphthalene	< 0.1	< 0.1	n.a.	n.d.	n.a.
Acenaphthylene	< 0.1	< 0.1	n.a.	n.d.	n.a.
Acenaphthene	< 0.1	< 0.1	n.a.	n.d.	n.a.
Fluorene	< 0.1	< 0.1	n.a.	0.2	n.a.
Phenanthrene	2.2	< 0.1	1.0	1.1	100
Anthracene	< 0.1	< 0.1	n.a.	0.3	n.a.
Fluoranthene	4.5	0.8	2.5	2.4	83
Pyrene	5.3	0.9	2.9	2.8	83
Benz[a]anthracene	1.8	< 0.1	0.8	1.0	100
Chrysene	2.1	< 0.1	0.9	1.1	100
Benzo[b]fluoranthene	2.1	< 0.1	1.0	1.5	100
Benzo[k]fluoranthene	2.6	< 0.1	1.2	2.8	100
Benzo[a]pyrene	3.1	< 0.1	1.4	1.2	100
Indeno[1,2,3-cd]pyrene	2.4	< 0.1	1.1	1.6	100
Dibenz[a,h]anthracene	< 0.1	< 0.1	n.a.	n.d.	n.a.
Benzo[g,h,i]perylene	2.0	< 0.1	0.9	1.5	100
Total (mg/kg)	28	2	14	17	93

NLU 54-SS-0010: Summary of PAH Analyses on Bulk Sediment Sample and Separated Fractions

Note:

Reported concentration values are average values of duplicate tests with relative standard deviation (RSD) ranging from 7~13% for all PAH compounds of interest.