

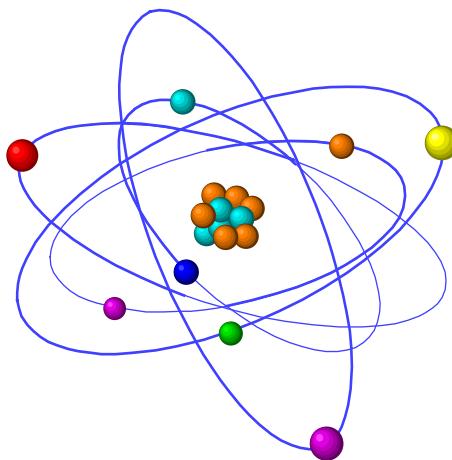
Strategic Oil & Gas Ltd.

Work Order-Ref #: 19130

Vapor Intrusion Assessment (VIA) Soils Outside Casing (AGM)

Strategic et al Cameron F-73

September 10 & 11, 2019



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FORENSIC SOLUTIONS FOR ENERGY CHALLENGES

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1.0 Executive Summary

On September 10-11, 2019, Strategic et al Cameron Hills F-73 was investigated for natural gas leakage in soils outside of casing. Total combustible gas (%LEL) field instruments were used to measure gas levels and types at soil test sites. Gas samples were collected, contained and preserved from soils outside casing (AGM) and for baseline comparison, background, background locations, ~30m away from the wellbore were also assessed.

Soils outside of casing immediately adjacent to of the wellbore were water saturated however, no gas bubbling was observed in standing water. A 113 site non-intrusive surface CH₄ scan was conducted in soils outside casing and at 3-background locations (W30m, S30m, and SE30m). Two sites (N1m-E4m and E3m) had elevated (18 and 11 ppm v/v, respectively) above background methane readings (2 ppm v/v). All other sites tested near the wellbore had CH₄ levels of 2 ppm v/v and were similar to the BKG sites.

Flux tests using GCHEM's Soil Vapor-Flux Chambers (SV-FC) conducted at the highest non-intrusive surface methane reading (N1m-E4m) and in background (BGK-SE30m) indicate very low combustible gas flow rates to surface (0.000035 and 0.000033 m³/day, respectively).

A total of 10 soil sites outside casing were assessed for gas leakage using an intrusive methodology where soil vapor test holes were augered into soils and Soil Vapor Probes (SVPs) were inserted into each test hole. Of the 10 soil vapor test sites outside casing, 9 sites (N3m 47 ppm v/v, N5m 13 ppm v/v, E1m 393 ppm v/v, E3m 67 ppm v/v, SE 1.5m 24,450 ppm v/v, S1m 16,900 ppm v/v, S3m 26,850 ppm v/v, SW1.5m 25,100 ppm v/v and W1m 11 ppm v/v) contained elevated, above background (SE30m 2 ppm v/v) methane contents. SOG selected 1 soil site (S3m) to measure chemical and $\delta^{13}\text{C}$ isotopic compositions to aid in classifying combustible gas contents. The 1-site contained high levels of CH₄ gas and low, similar to background levels of associated ΣC_2+ gas. $\delta^{13}\text{C}$ CH₄ and $\delta^{13}\text{C}$ CO₂ values were depleted when compared to thermogenic CH₄ gas and suggest that elevated %LEL values and associated CH₄ gas measured at the soil site is biogenic in origin where CH₄ is being generated via CO₂ reduction or fermentation pathways and then further altered via bacterial oxidation processes. ΣC_2+ gases in soils outside casing at S3m are the result of natural movement of light hydrocarbon gases from reservoirs at depth, upward through subsurface fractures and micro-fractures to surface. This is a naturally occurring process in all hydrocarbon sedimentary basins in the world.

With information available to date, soil vapor tests sites S3m would be classified as 'Biogenic-Naturally Occurring CH₄-Non-Impacted'.

2.0 Vapor Intrusion Assessment Summary

Operating Company: Strategic Oil & Gas Ltd.
Well Name: Strategic et al Cameron F-73
UWI: 300F7360117150

License Number: 001992
Test Date September 10-11, 2019
GCHEM Project Number 19130

2.1 Production Casing Test Summary Table

Combustible Gas (CH₄) ([%LEL])	nm		
Hydrogen Sulphide (H₂S) Gas (ppm v/v)	nm		
PC Flow Rate (m³/day)	nm		
P-T Date Logger Installed	nm		
P-T Data Logger Removed	nm		
P-T Data Logger Test Duration	nm		
MAX Pressure (kPa)	nm		
Gas Spls. Collection-Measurement	Total Collected	Analysis Requested*	Classification**
PC Samples (Total)	0		
PC Combustible Gas Class. Level-1 (Chemical)	NA	NA	NA
PC Combustible Gas Class. Level-2 (¹³C)		NA	NA
PC Combustible Gas Class. Level-3 (⁸D)		NA	NA
PC Combustible Gas Class. Level-4 (¹⁴C)		NA	NA

2.2 Surface Casing Vent Flow (SCVF) Test Summary Table

SCV Ten-Minute Bubble Test Result	nm		
SCV Flow Rate (m³/day)	nm		
SCVF Pressure-Temp Logger Installed	nm		
SCV Pressure-Temp Data Logger Removed	nm		
SCV Shut-In Time (hrs)	nm		
SCV MAX-Recorded Build Up Pressure (kPa)	nm		
SCV Stabilized Build-up Pressure (kPa):	nm		
SCV Stabilized Build-up Time (hours)	nm		
SCV Standpipe Max CH₄ Content (% LEL):	nm		
SCV Standpipe Max H₂S Content	nm		
SCV Gas Spls. Collection-Measurement	Total Collected	Analysis Requested*	Classification**
SCV Samples (Total)	0		
SCV Combustible Gas Class. Level-1 (Chemical)	NA	NA	NA
SCV Combustible Gas Class. Level-2 (¹³C)		NA	NA
SCV Combustible Gas Class. Level-3 (⁸D)		NA	NA
SCV Combustible Gas Class. Level-4 (¹⁴C)		NA	NA

2.3 Soil Gas Migration-Vapor Intrusion Assessment: Soils Outside Casing (AGM) Summary

A) Non-Intrusive CH₄ Surface Soil Scan (PMD) (Figure-1 and Table 1)

Well Casing Surface CH ₄ Test Sites	113
MAX Surface CH ₄ Reading	18
MAX H ₂ S Well Soil Reading (ppm v/v)	0
Number of Background Sites	3
MAX Background CH ₄ (ppm v/v)	2
Max H ₂ S BKG Soil Reading (ppm v/v)	0
Surface CH ₄ -PMD Gas Classification	

B) Non-Intrusive Surface Enclosed Soil Vapor FLUX Chamber Test

Surface SV-FC CH ₄ Test Sites	N1m-E4m, BKG SE30m		
MAX SV-FC CH ₄ Reading	18		
SV-FC Gas Spls. Collection-Measurement	Total Collected	Analysis Requested*	Test Site
SV-FC Samples (Total)	2		
SV-FCs and Sites Requested for Level-1 Analysis		0	N/A
Combustible Gas Classification Level-1 (Chem.)			N/A
SV-FCs and Sites Requested for Level-2 Analysis		0	N/A
Combustible Gas Classification Level-2 ($\delta^{13}\text{C}$)			N/A
SV-FCs and Sites Requested for Level-3 Analysis		0	N/A
Combustible Gas Classification Level-3 (δD)			N/A
SV-FCs and Sites Requested for Level-4 Analysis		0	N/A
Combustible Gas Classification Level-4 ($\delta^{14}\text{C}$)			N/A

C) Intrusive Auger Test Holes with Soil Vapor Sampling or Soil Vapor Monitoring Probes

Number Soil Vapor Probe (SVP) Test Sites	10		
MAX SVP CH ₄ Reading (%LEL)	53.7		
Max H ₂ S SVP Field Reading (ppm v/v)	0		
Number SVP BKG Test Sites	3		
MAX SVP CH ₄ BKG Test Sites (ppm v/v)	2		
SVPs Gas Spl. Collection & Measurement	Total Collected	Analysis Requested*	Test Site
Soil Vapor Probes (SVPs) AGM (Total)	3		
SV-FCs and Sites Requested for Level-1 Analysis		1	S3m
Combustible Gas Classification Level-1 (Chem.)			Biogenic-Non-Impacted
SV-FCs and Sites Requested for Level-2 Analysis		1	S3m
Combustible Gas Classification Level-2 ($\delta^{13}\text{C}$)			Biogenic-Non-Impacted
SV-FCs and Sites Requested for Level-3 Analysis		0	NA
Combustible Gas Classification Level-3 (δD)			NA
SV-FCs and Sites Requested for Level-4 Analysis		0	NA
Combustible Gas Classification Level-4 ($\delta^{14}\text{C}$)			NA



BKG Gas Spl. Collection-Measurement	Total Collected	Analysis Requested*	Test Site
BKG Soil Vapor Probe (SVPs) (Total)	3		
SV-FCs and Sites Requested for Level-1 Analysis		1	BKG SE30m
Combustible Gas Classification Level-1 (Chem.)			Biogenic-Naturally Occurring-Baseline
SV-FCs and Sites Requested for Level-2 Analysis		0	N/A
Combustible Gas Classification Level-2 ($\delta^{13}\text{C}$)			NA
SV-FCs and Sites Requested for Level-3 Analysis		0	N/A
Combustible Gas Classification Level-3 (δD)			NA
SV-FCs and Sites Requested for Level-4 Analysis		0	N/A
Combustible Gas Classification Level-4 ($\delta^{14}\text{C}$)			NA

* *Sample selection for chemical and isotopic analysis (geochemical analytical suite) selected by client/operator.*

2.4 Interpreted Source of Natural Gas Found at/near Surface: measured depth from KB of the well (Figures 3 to 6).

Sample Point	Geologic Formation	Depth Range	Source Depth
SVP S3m	Near Surface Soil Respiration		Biogenic CH ₄ , Non-Impacted, Baseline



3.0 Background of Vapor Intrusion Assessments (VIA) at Resource Wells & Tracing Gas Contents in the Environment using Energy Forensics

Undesired natural gas leakage from depth to surface at resource wells is becoming increasingly recognized and is a significant financial burden to the resource industry. When high levels of natural gas are found in the surface casing vent it is termed surface casing vent flow (SCVF) and when found in soils outside casing it is termed active gas migration (AGM). Identifying the source of leaking gas, maintaining zonal isolation and eliminating gas leakage to surface has proven to be a challenging task. Industry success rates using conventional gas leakage identification tools (e.g. noise, temperature, cement bond-integrity, ultra-sonic imaging logs, etc.) to eliminate surface gas migration in the first attempt is approximately 15% to 20%. Since 1997, through collaboration with industry, government regulators and academic institutions, GCHEM Ltd. has developed ‘Energy-Forensics’ and has obtained extensive expertise in field testing, gas sampling and preservation, analytical and interpretational techniques to pinpoint the geologic source of natural gases at resource wells.

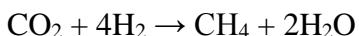
It is important to note that detection of elevated combustible gases at surface does not always mean the well is impacted with deep sourced natural gas (thermogenic). Accurate gas characterization at well sites is critical as elevated CH₄ (%LEL) contents measured at or near surface may not indicate it is leaking or impacted (false-positive) but rather the combustible gases present are the result of biogenic activity or hydrocarbon contamination (or a combination of).

Thermogenic hydrocarbon gases have unique chemical and isotopic signatures based on many variables including the starting organic material they are produced from, the chemical processes from organic origin to current form, interaction with surrounding formation rock and fluids, and effects from migrating from origin to current trap. For example, molecular and isotopic composition ($\delta^{13}\text{C}$ and $\delta^2\text{H}$) of a low temperature, shallow sourced natural gas is significantly different with respect to those of a high temperature deep sourced natural gas. This principle allows the geologic source of leaking natural gas at a wellbore to be determined.

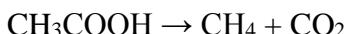
3.1 Biogenically Derived Methane Gas

As a normal part of soil respiration, methane may be generated via two biotic pathways (depending on substrate availability):

CO₂ reduction reaction



Fermentation Process



Biogenic methane gas may be further oxidized by bacteria. Oxidizing bacteria in soils preferentially consume ¹²C over ¹³C resulting the residual gas enriched in ¹³C (i.e. $\delta^{13}\text{C}$ values become less negative) with respect to the biogenic gas (isotope enriching effect). Therefore, biogenic oxidation may provide the false signature of a ‘mixture of biogenic and thermogenic methane’ or ‘thermogenic’ gas (GCHEM in prep). Biogenic CH₄ is generally prevalent in landfill or swamp gas.

3.2 Thermogenic Methane Gas

Methane gas can be generated by abiotic processes such as the thermo-degradation of organic matter at high pressure and temperature (thermogenesis). During thermogenic CH₄ generation, pending organic matter content, pressure and temperature, associated C₂+ gases may also be formed. Thermogenic CH₄ and C₂+ gases contain enriched $\delta^{13}\text{C}$ and δD values pending gas maturity, mixing and alteration and torturous pathway from source to trap.

3.3 Classification, Characterization and Geological Origins of Combustible Gases in the Environment.

Combustible gases in soils outside casing maybe classified and characterized (biogenic, thermogenic or mixed) using chemical, carbon and hydrogen and isotopic measurements and ¹⁴C concentrations. Leaking thermogenic natural gas in soils outside casing is easier to scientifically prove than biogenic methane sources. Elevated %LEL measured in AGM (on location) maybe the result of naturally occurring biogenic processes, anthropogenic leaking thermogenic natural gases and mixtures of both. A systematic 4-level approach can be used to determine the origins (biogenic-thermogenic or mixed) combustible gas contents and include:

1) Level-1 Characterization**High Resolution Compositional-Chemical Measurements.**

Permanent, inert and CH₄ to n-C₅H₁₂ & C₆+

See NGGC-1 CH₄ vs ΣC₂+ (Szatkowski et al 2000 & 2001).

See NGGC-2 C₂H₆ vs. C₆+ (Szatkowski et al 2000 & 2001).

2) Level-2 Characterization**Stable Carbon Isotope Measurements (δ¹³C).**

δ¹³C CH₄ to n-C₅H₁₂ & CO₂ (pending concentrations-gas levels).

See NGGC-3 CH₄/ΣC₂+ vs. δ¹³C CH₄ (Bernard 1978).

See NGGC-4 δ¹³C CO₂ vs. δ¹³C CH₄ (Whiticar 1993).

3) Level-3 Characterization**Hydrogen in Methane (δD).**

δD CH₄ to dD C₄H₁₂ (pending concentrations-gas levels).

See NGGC-5 δ¹³C CH₄ vs δD CH₄ (Coleman 1993).

4) Level-4 Characterization**¹⁴C pMC concentrations (radioactive ½ life of 5750 yr).**

Pending concentrations-gas levels.

¹⁴C reveals the age of the organic matter source from which CH₄ was generated but not the time of methanogenesis.

To determine the geological origins of leaking thermogenic natural gas contents, a series of plots developed by GCHEM Ltd are used and include.

- | | |
|---|--|
| 1) Chemical & Isotopic Gas Field Diagram | C ₂ H ₆ /ΣC ₃ + vs δ ¹³ C C ₂ H ₆ (Szatkowski et al 2000, 2001). |
| 2) Isotopic Gas Field Diagram | δ ¹³ C C ₂ H ₆ vs. δ ¹³ C C ₃ H ₈ (Szatkowski et al 2000, 2001). |
| 3) Modified Chung Plot | δ ¹³ C vs 1/n (carbon & hydrogen number) (Chung 1988, and GCHEM Ltd. Unpublished). |

Additional chemical and stable carbon and hydrogen isotopic plots have been developed to aid in determining the geological origins of natural gas found in the environment however, GCHEM has not published these novel and new correlations and relationships and they will not be shown or discussed in detail at this time (GCHEM Unpublished Internal Research).

4.0 Methods and Results

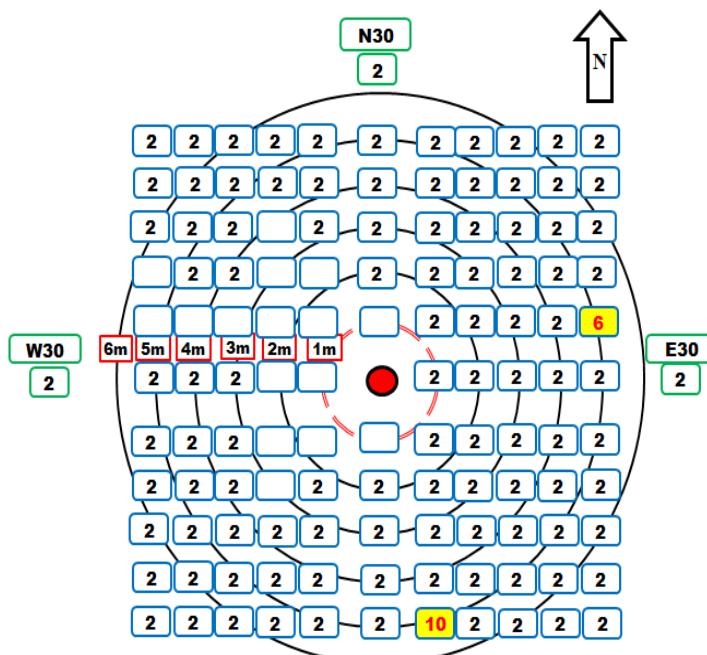
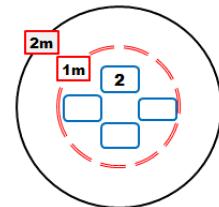
4.1 Field Assessment Methods and Results

4.1.1 Non-Intrusive Vapour Intrusion Assessment

On September 10, 2019, GCHEM conducted a surface soil methane scan using a Sensit PMD (Figure 1). CH₄ readings were measured at 113 locations on a grid pattern (1m x 1m) covering approximately an 10m x 10m square area around the marked wellbore. Soils outside casing were wet (standing water in some areas) however, no gas bubbling in standing water was observed while conducting the VIA in soils outside of casing.

To establish background surface CH₄ gas levels a distance away from the well bore, three locations (30m west, 30m south, and 30m southeast of the wellbore) were also assessed. To enhance results of the surface methane scan and reduce potential effects from industrial contamination, at each test site, an atmospheric CH₄ gas level was recorded, the PMD gas sampling wand was coupled to surface soils and the CH₄ level was recorded for that specific test site. Atmospheric CH₄ level was subtracted from the CH₄ level measured after ground coupling to derive a surface soil CH₄ level at that point of the grid.

Two sites (N1m-E4m and E3m) had elevated (18 and 11 ppm v/v, respectively) above background methane readings (2 ppm v/v). All other sites tested near the wellbore had CH₄ levels of 2 ppm v/v and were similar to the BKG sites.

Figure 1. AGM Non-Intrusive Surface PMD

Figure-1A. NON-Intrusive CH₄ Surface Scan Well Casing Detail VIEW


Seeded grass surrounding the well casing is of similar stand-growth to surrounding lease vegetation. No stressed dead spots or discoloration was observed.

Table 1. AGM Non-Intrusive Surface PMD

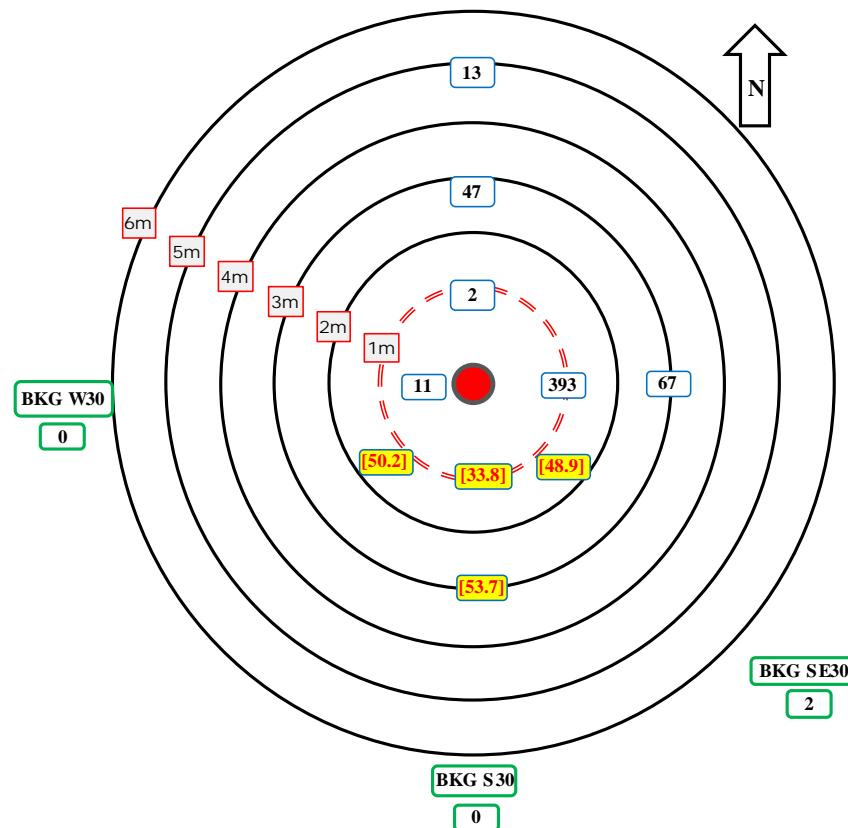
Test Site (m)	WELL CASING (AGM) Non-Intrusive Surface PMD (CH ₄)		Test Site (m)	PMD (CH ₄)		Test Site (m)	PMD (CH ₄)		Test Site (m)	Soil Scan PMD CH ₄	
	(ppm v/v)	(% Vol)		(ppm v/v)	(% Vol)		(ppm v/v)	(% Vol)		(ppm v/v)	(% Vol)
N.5	2		E.5			S.5			W.5		
N1	2		E1	2		S1	2		W1		
N2	2		E2	2		S2	2		W2	2	
N3	2		E3	11		S3	2		W3	2	
N4	2		E4	2		S4	2		W4	2	
N5	2		E5	2		S5	2		W5	2	
N5-E1	2		E5-S 1	2		S5-W1	2		W5-N1	2	
N4-E1	2		E5-S 2	2		S4-W1	2		W5-N2	2	
N3-E1	2		E5-S 3	2		S3-W1	2		W5-N3	2	
N2-E1	2		E5-S 4	2		S2-W1	2		W5-N4	2	
N1-E1	2		E5-S 5			S1-W1	2		W5-N5	2	
N1-E2	2		E4-S 5	2		S1-W2	2		W4-N5	2	
N2-E2	2		E4-S 4	2		S2-W2	2		W4-N4	2	
N3-E2	2		E4-S 3	2		S3-W2	2		W4-N3	2	
N4-E2	2		E4-S 2	2		S4-W2	2		W4-N2		
N5-E2	2		E4-S 1	2		S5-W2	2		W4-N1		
N5-E3	2		E3-S 1	2		S5-W3	2		W3-N1		
N4-E3	2		E3-S 2	2		S4-W3	2		W3-N2	2	
N3-E3	2		E3-S 3	2		S3-W3	2		W3-N3	2	
N2-E3	2		E3-S 4	2		S2-W3	2		W3-N4	2	
N1-E3	2		E3-S 5	2		S1-W3			W3-N5	2	
N1-E4	18		E2-S 5	2		S1-W4	2		W2-N5	2	
N2-E4	2		E2-S 4	2		S2-W4	2		W2-N4	2	
N3-E4	2		E2-S 3	2		S3-W4	2		W2-N3	2	
N4-E4	2		E2-S 2	2		S4-W4	2		W2-N2	2	
N5-E4	2		E2-S 1	2		S5-W4	2		W2-N1	2	
N5-E5	2		E1-S 1	2		S5-W5	2		W1-N1	2	
N4-E5	2		E1-S 2	2		S4-W5			W1-N2	2	
N3-E5	2		E1-S 3	2		S3-W5	2		W1-N3	2	
N2-E5	2		E1-S 4	2		S2-W5	2		W1-N4	2	
N1-E5	2		E1-S 5	2		S1-W5	2		W1-N5	2	

BACKGROUND Non-Intrusive Surface PMD (CH₄) Soil Scan

Test Site (m)	PMD CH ₄ (ppm v/v)	(%)	Test Site (m)	PMD CH ₄ (ppm v/v)	(%)	Test Site (m)	PMD CH ₄ (ppm v/v)	(%)	Test Site (m)	PMD CH ₄ (ppm v/v)	(%)
SE30	2		S30	2		S10	2		W30	2	

4.1.2 Intrusive Vapor Intrusion Assessment

A total of 10 soil sites outside casing were assessed for gas leakage using an intrusive methodology where soil vapor test holes were augered into soils and Soil Vapor Probes (SVPs) were inserted into each test hole. The SVPs were allowed to stabilize for approximately 30-minutes prior to combustible gas content measurement with the PMD (Figure 2). Soils were water saturated at depth below 1.5m thus a full intrusive 16-auger test pattern could not be conducted. Of the 10 soil vapor test sites, 9 sites (N3m 47 ppm v/v, N5m 13 ppm v/v, E1m 393 ppm v/v, E3m 67 ppm v/v, SE 1.5m 24,450 ppm v/v, S1m 16,900 ppm v/v, S3m 26,850 ppm v/v, SW1.5m 25,100 ppm v/v and W1m 11 ppm v/v) contained elevated, above background (SE30m 2 ppm v/v) methane contents.

Figure 2. AGM Intrusive SVPs


Seeded grass surrounding the well casing is of similar stand-growth to surrounding lease vegetation. No stressed dead spots or discoloration was observed.

Table 2. AGM Intrusive SVPs

Intrusive AGM - Hand Auger-Test Hole-Install Soil Vapor Probes (SVPs) ATM-Isolated						
Test Site (m)	Soil Vapor Probes		Soil Parameters		Gas Sample (Y-N)	Site Assessment Comments
	IR-CH ₄ (ppm v/v)	H ₂ S (ppm v/v)	Type	Moist. HC-CONT (1-5)		
N1	2	<1.0	Si	5	No	
N3	47	<1.0	Si	5	No	
N5	13	<1.0	Si	5	No	
E1	393	<1.0	Si	5	No	
E3	67	<1.0	Si	5	No	
E5						
SE1.5	24450	[48.9]	<1.0	Si	5	No
S1	16900	[33.8]	<1.0	Si	5	No
S3	26850	[53.7]	<1.0	Si	5	No
S5						
SW1.5	25100	[50.2]	<1.0	Si	5	No
W0.5	11	<1.0	Si	5	No	
W1						
W3						
W5						
Final SVP-Watered Out						
Test Site (m)	Soil Vapor Probes		Soil Parameters		Gas Sample (Y-N)	Site Assessment Comments
	IR-CH ₄ (ppm v/v)	H ₂ S (ppm v/v)	Type	Moist. HC-CONT (1-5)		
BKG SE30	2	<1.0	Si	5	No	Yes
BKG W30	0	<1.0	Si	5	No	Yes
BKG S30	0	<1.0	Si	5	No	Yes

4.1.3 Soil Vapor Flux Measurements

Soil Vapor flux measurements can be conducted in soils to establish the rate and volume of gas leakage at surface. The soil Vapor flux methodology utilizes an enclosed chamber (known internal volume and surface soil area) with three gas ports: gas-in, gas-out and a pressure release valve. Gases are cycled from the gas-out port to a PMD and re-injected or cycled back into the flux chamber. The atmospheric pressure release valve allows leaking gas from soils to enter the chamber and displace atmospheric gas contents within the chamber.

Pristine, naturally occurring gas venting from soils as a result of natural movement of light hydrocarbons from reservoirs at depth, upward through subsurface fractures or micro-fractures to surface combined with soil respiration processes is a naturally occurring process prevalent in all sedimentary basins (i.e. hydrocarbon surface seeps). These soil gases are usually comprised of low, but variable, levels of CH₄ and CO₂ with low-to-trace levels of associated C₂₊ thermogenic natural gases that cannot be generated by bacterial processes in great quantities. Soils influenced by anthropogenic process (i.e. natural gas leakage at a wellbore from natural gas reservoirs at depth, upwards through compromised cement sheaths securing production casing to formation rock to surface) usually contain highly elevated, above background levels of CH₄ (thermogenic, biogenic and/or mixtures) and associated C₂₊ thermogenic gases.

CH₄ gas contents in the flux chamber were monitored and data logged using a PMD. Soil gas flux volumes and rates in soils can be calculated either volumetrically or gravimetrically considering the following relationship:

$$\text{Flux (F)} = (\text{dC}/\text{dt}) * (\text{volume}) / (\text{area})$$

Where:

C = concentration

t = time

dC/dt = change in concentration with time (the slope of a *concentration versus elapsed time* plot).

The volumetric flux is calculated from ppm v/v units (10⁻⁶ m³ light alkanes/ m³ air) and the gravimetric flux is calculated by converting ppm v/v to g/m³ of air using the ideal gas law (PV = nRT) with P = 1 atmosphere and T = 25°C.

On September 10, 2019 the highest methane reading observed during the non-intrusive surface scans was at N1m-E4m (18 ppm v/v methane). A GCHEM soil vapour-flux chamber (SV-FC) was installed at this location and a 27-minute flux was conducted (Figure-3, Table-3). At the start of the flux the combustible gas reading was 44 ppm v/v methane and did not significantly change throughout the test. A gas flow rate of 0.000035 m³/day was calculated (Table 5). Gas samples were collected from the flux chamber at the conclusion of the test (27 minutes).

A second flux test was performed at BKG-SE30m (Figure-3, Table-4). At the start of the flux the combustible gas reading was 0 ppm v/v methane and increased to about 5 ppm v/v methane during the 31-minute test. A gas flow rate of 0.000033 m³/day was calculated (Table 6). Gas samples were collected from the flux chamber at the conclusion of the test (31 minutes).

Strategic et al Cameron F-73 Soil Vapor Flux Field CH ₄ Test Time & Levels Site N1mE4m											
Test-Time	PPM	%LEL	%V/V	Test-Time	PPM	%LEL	%V/V	Test-Time	PPM	%LEL	%V/V
17:27:00	44	0	0	17:38:00	46	0	0	17:49:00	40	0	0
17:27:30	46	0	0	17:38:30	43	0	0	17:49:30	42	0	0
17:28:00	46	0	0	17:39:00	44	0	0	17:50:00	43	0	0
17:28:30	45	0	0	17:39:30	45	0	0	17:50:30	44	0	0
17:29:00	47	0	0	17:40:00	43	0	0	17:51:00	42	0	0
17:29:30	46	0	0	17:40:30	43	0	0	17:51:30	42	0	0
17:30:00	46	0	0	17:41:00	42	0	0	17:52:00	42	0	0
17:30:30	46	0	0	17:41:30	43	0	0	17:52:30	44	0	0
17:31:00	45	0	0	17:42:00	45	0	0	17:53:00	40	0	0
17:31:30	45	0	0	17:42:30	42	0	0	17:53:30	43	0	0
17:32:00	44	0	0	17:43:00	44	0	0	17:54:00	41	0	0
17:32:30	46	0	0	17:43:30	42	0	0				
17:33:00	47	0	0	17:44:00	44	0	0				
17:33:30	45	0	0	17:44:30	45	0	0				
17:34:00	45	0	0	17:45:00	43	0	0				
17:34:30	45	0	0	17:45:30	43	0	0				
17:35:00	45	0	0	17:46:00	44	0	0				
17:35:30	44	0	0	17:46:30	41	0	0				
17:36:00	45	0	0	17:47:00	43	0	0				
17:36:30	46	0	0	17:47:30	42	0	0				
17:37:00	46	0	0	17:48:00	42	0	0				
17:37:30	46	0	0	17:48:30	43	0	0				

Table-3. CH₄ Gas Levels vs. Test measured by the PMD in the FLUX Chamber at N1m-E4m at Strategic et al Cameron F-73.

Strategic et al Cameron F-73 Soil Vapor Flux Field CH ₄ Test Time & Levels BKG 30m SE											
Test-Time	PPM	%LEL	%V/V	Test-Time	PPM	%LEL	%V/V	Test-Time	PPM	%LEL	%V/V
17:27:38	0	0	0	17:38:38	3	0	0	17:49:38	5	0	0
17:28:08	0	0	0	17:39:08	2	0	0	17:50:08	6	0	0
17:28:38	0	0	0	17:39:38	4	0	0	17:50:38	5	0	0
17:29:08	0	0	0	17:40:08	2	0	0	17:51:08	5	0	0
17:29:38	0	0	0	17:40:38	3	0	0	17:51:38	5	0	0
17:30:08	0	0	0	17:41:08	3	0	0	17:52:08	5	0	0
17:30:38	0	0	0	17:41:38	4	0	0	17:52:38	5	0	0
17:31:08	0	0	0	17:42:08	4	0	0	17:53:08	5	0	0
17:31:38	0	0	0	17:42:38	4	0	0	17:53:38	6	0	0
17:32:08	0	0	0	17:43:08	4	0	0	17:54:08	5	0	0
17:32:38	0	0	0	17:43:38	4	0	0	17:54:38	5	0	0
17:33:08	0	0	0	17:44:08	4	0	0	17:55:08	5	0	0
17:33:38	0	0	0	17:44:38	4	0	0	17:55:38	5	0	0
17:34:08	0	0	0	17:45:08	6	0	0	17:56:08	6	0	0
17:34:38	0	0	0	17:45:38	4	0	0	17:56:38	5	0	0
17:35:08	0	0	0	17:46:08	5	0	0	17:57:08	5	0	0
17:35:38	0	0	0	17:46:38	5	0	0	17:57:38	6	0	0
17:36:08	1	0	0	17:47:08	5	0	0	17:58:08	5	0	0
17:36:38	2	0	0	17:47:38	5	0	0	17:58:38	5	0	0
17:37:08	3	0	0	17:48:08	4	0	0				
17:37:38	4	0	0	17:48:38	5	0	0				
17:38:08	2	0	0	17:49:08	6	0	0				

Table-4. CH₄ Gas Levels vs. Test Time measured by the PMD in the FLUX Chamber at BKG SE 30m from at Strategic et al Cameron F-73.

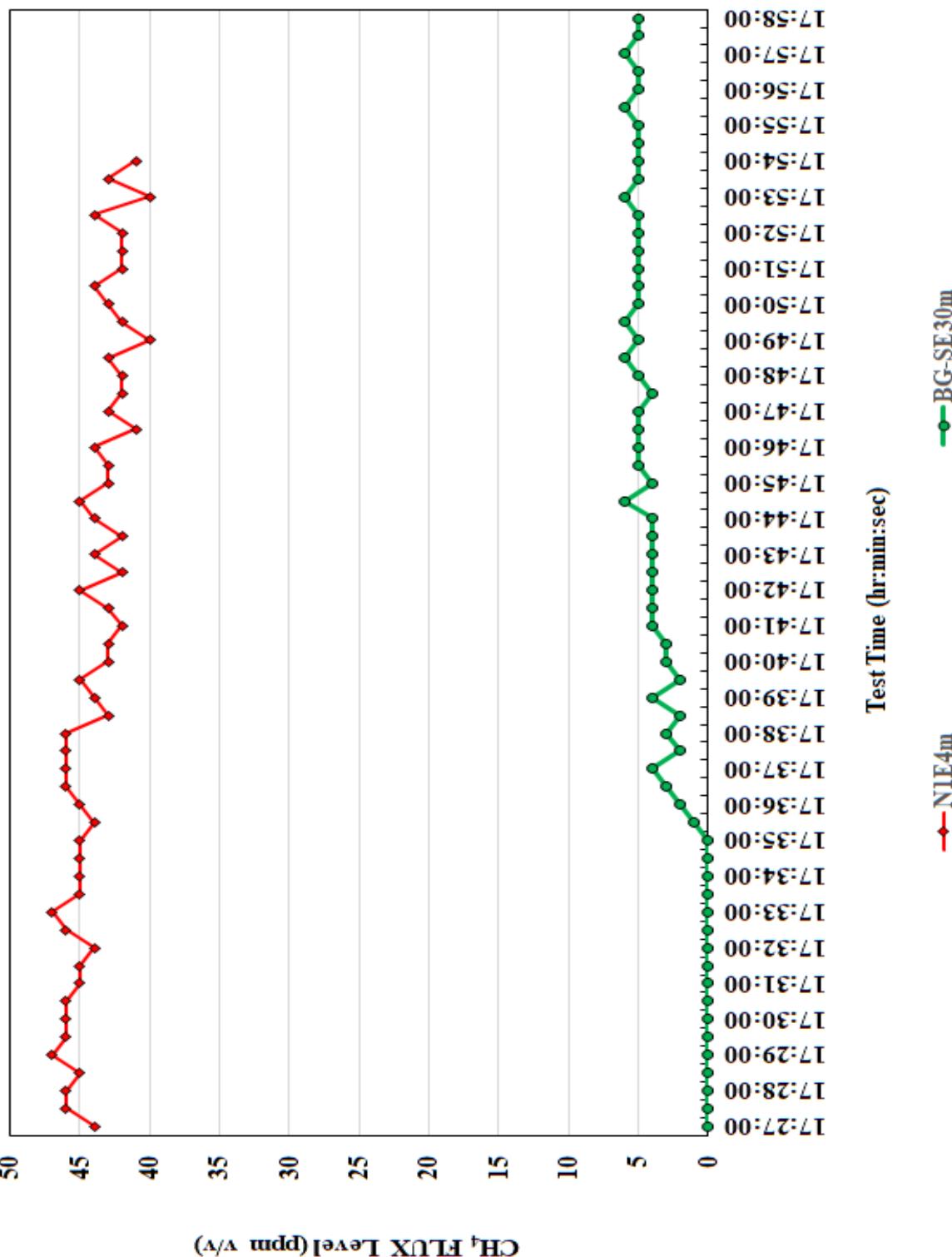
Strategic et al Cameron F-73
Soil Vapor Flux Chambers(SV-FC)
September 10-2019

Figure-3. CH₄ Gas Levels vs. Test Time measured by the PMD in the FLUX Chamber at anomalous AGM Site N1m-E4m and BKG SE30m from the well head at Strategic et al Cameron F-73.

Field PMD CH ₄ Values			
Venting Gas Volume Calculation Type	Gas Component	Volumetric FLUX (m ³ /m ² /day)	Gravimetric FLUX (g/m ² /day)
CH ₄ Gas FLUX Volume	Methane (CH ₄)	0.000035	0.021017068
Laboratory Chemical Compositions			
	Gas Component	Volumetric FLUX (m ³ /m ² /day)	Gravimetric FLUX (g/m ² /day)
Speciated LHG & CO ₂ Gas FLUX Volume	Methane (CH ₄) Ethane (C ₂ H ₆) Propane (C ₃ H ₈) n-Butane (n-C ₄ H ₁₀) Carbon Dioxide (CO ₂)		

Table-5. Calculated venting CH₄ (methane) gas FLUX & Speciated FLUX Rate-Volume measured at AGM Site N1m-E4m at Strategic et al Cameron F-73.

Field PMD CH ₄ Values			
Venting Gas Volume Calculation Type	Gas Component	Volumetric FLUX (m ³ /m ² /day)	Gravimetric FLUX (g/m ² /day)
CH ₄ Gas FLUX Volume	Methane (CH ₄)	0.000033	0.021017068
Laboratory Chemical Compositions			
	Gas Component	Volumetric FLUX (m ³ /m ² /day)	Gravimetric FLUX (g/m ² /day)
Speciated LHG & CO ₂ Gas FLUX Volume	Methane (CH ₄) Ethane (C ₂ H ₆) Propane (C ₃ H ₈) n-Butane (n-C ₄ H ₁₀) Carbon Dioxide (CO ₂)		

Table-6. Calculated venting CH₄ (methane) gas FLUX & Speciated FLUX Rate-Volume measured at BKG SE30m at Strategic et al Cameron F-73.

4.2 Analytical Methods

- a. High Resolution Compositional Analysis (HRCA).
 - i. He, H₂, O₂, N₂, CO₂, CH₄ to n-C₅H₁₂ & C₆+
- b. Stable Carbon ($\delta^{13}\text{C}$) and Hydrogen (δD) Isotopic Analysis.
 - i. $\delta^{13}\text{C}$ CH₄ to n-C₅H₁₂ and CO₂, and δD CH₄ to n-C₅H₁₂

Compositional (molecular) analyses were conducted at GCHEM's Analytical Laboratory using Hewlett Packard 5890 and Agilent 7890 Gas Chromatographs (GCs) configured for low (ppb v/v to ppm v/v) to high (vol. %) level detection of light alkane/alkene gases and atmospheric gas components. Chemical analysis of gases measured, and analytical error are shown in Table-1.

Stable carbon ($\delta^{13}\text{C}$) isotope ratios of light hydrocarbon gases (LHG) and carbon dioxide and hydrogen isotope ratios (δD) of LHG were also measured at GCHEM's Analytical Laboratory on a Thermo-Scientific MAT-253 Gas Chromatograph-Combustion-Continuous Flow-Isotope Ratio Mass Spectrometer (GC-C-CF-IRMS). Carbon isotope ratios are reported in delta (δ) notation and per mil (‰, parts per thousand) with respect to VPDB (Vienna Pee Dee Belemnite). Hydrogen isotope ratios are reported in delta (δ) notation and per mil (‰) with respect to VSMOW (Vienna Standard Mean Ocean Water).

Gas Component (ppmv)	Molecular Formula	Analytical Error (%)	Stable Carbon Isotopic Composition ($\delta^{13}\text{C}$)	Analytical Error (‰ VPDB)	Hydrogen Isotopic Composition (δD)	Analytical Error (‰ VSMOW)
Hydrogen	H ₂	±7%	-	-	$\delta\text{D H}_2$	±10
Helium	He	±7%	-	-	-	-
Nitrogen	N ₂	±7%	-	-	-	-
Oxygen	O ₂	±7%	-	-	-	-
Carbon Dioxide	CO ₂	±7%	$\delta^{13}\text{C CO}_2$	±0.2	-	-
Hydrogen Sulphide	H ₂ S	±7%	-	-	-	-
Methyl Mercaptan	CH ₄ S	±7%	-	-	-	-
Ethyl Mercaptan	C ₂ H ₆ S	±7%	-	-	-	-
Thiophene	C ₄ H ₄ S	±7%	-	-	-	-
Dimethyl Disulfide	C ₂ H ₆ S ₂	±7%	-	-	-	-
Methane	CH ₄	±7%	$\delta^{13}\text{C CH}_4$	±0.1	$\delta\text{D CH}_4$	±10
Ethane	C ₂ H ₆	±7%	$\delta^{13}\text{C C}_2\text{H}_6$	±0.2	$\delta\text{D C}_2\text{H}_6$	±10
Ethene	C ₂ H ₄	±7%	$\delta^{13}\text{C C}_2\text{H}_4$	±0.2	$\delta\text{D C}_2\text{H}_4$	±10
Propane	C ₃ H ₈	±7%	$\delta^{13}\text{C C}_3\text{H}_8$	±0.2	$\delta\text{D C}_3\text{H}_8$	±10
Propene	C ₃ H ₆	±7%	$\delta^{13}\text{C C}_3\text{H}_6$	±0.2	$\delta\text{D C}_3\text{H}_6$	±10
iso-Butane	i-C ₄ H ₁₀	±7%	$\delta^{13}\text{C i-C}_4\text{H}_{10}$	±0.2	$\delta\text{D i-C}_4\text{H}_{10}$	±10
normal-Butane	n-C ₄ H ₁₀	±7%	$\delta^{13}\text{C n-C}_4\text{H}_{10}$	±0.2	$\delta\text{D n-C}_4\text{H}_{10}$	±10
iso-Pentane	i-C ₅ H ₁₂	±7%	$\delta^{13}\text{C i-C}_5\text{H}_{12}$	±0.2	$\delta\text{D i-C}_5\text{H}_{12}$	±10
normal-Pentane	n-C ₅ H ₁₂	±7%	$\delta^{13}\text{C n-C}_5\text{H}_{12}$	±0.2	$\delta\text{D n-C}_5\text{H}_{12}$	±10
Hexane and higher	C ₆ +	±7%	-	-	-	-

Table 7. Gas components, isotopic compositions measured and the analytical error of the measurements at GCHEM's Analytical Laboratory.



5.0 Geochemical Measurements-Laboratory Results.

As part of this VIA (SCV-AGM), a total of 4 gas samples were collected, contained and preserved from the following locations or sample points: SVPs-soils outside casing (S1m, S3m, SE1.5m and N1-E4 Chamber) and 4-BKG locations (W30m, SW30m, SE30m and E30m).

At the request of the Strategic Oil and Gas, chemical and $\delta^{13}\text{C}$ isotopic compositions were measured for gases obtained from one (S3m) of the SVPs that contained elevated, above background, levels of combustible gases and one BKG SVP (SE30m). High Resolution chemical and $\delta^{13}\text{C}$ isotopic compositions were measured at GCHEM's Forensic Lab and are provided in Table 8.

5.1 Gases Obtained from Soil Vapor Probes (SVPs).

Gases measured in the SVPs in soils near the well bore (S3m) contain above atmospheric levels of CO_2 (10,811 ppm v/v). Methane gas was elevated (2368 ppm v/v) when compared to background level measured at BKG SE30m (2.48 ppm v/v) (Table 8 and Figure 5). C_2+ gas levels in SVP S3m were low (3.14 ppm v/v) and were similar to background levels (<0.01 ppm v/v). High methane with low, associated C_2+ thermogenic gases suggests a biogenic or biotic source via CO_2 reduction or fermentation reactions for methane gas. This gas has then likely undergone secondary bacterial oxidation processes (Figure 6, and Figure 7). C_6+ gas contents at SVP site S3m were low (0.26 ppm v/v respectively) and suggest hydrocarbon contamination was not present at SVP test sites (Figure 5).

C_2+ gas levels at SVP site S3m were too low to measure $\delta^{13}\text{C}$ isotopic compositions. Sufficient levels of CH_4 and CO_2 were available for $\delta^{13}\text{C}$ at site S3m. $\delta^{13}\text{C}$ CH_4 and $\delta^{13}\text{C}$ CO_2 at SVP S3m was -50.39 and -11.93‰ VPDB respectively (Table 8). These values are consistent with gases originating from a biogenic source that have undergone secondary bacterial oxidation processes (Figures 6 and 7).

Table 8. High resolution molecular and stable carbon isotopic compositions of gas samples collected as part of the VIA at Strategic et al Cameron F-73. Hydrogen isotopic compositions were not measured at the request of SOG.

Gas Component	Sample Point	S3	BKG SE
	Date Collected	Sept. 10-19 (ppm v/v)	Sept. 10-19 (ppm v/v)
Hydrogen		3.40	3.13
Helium		2.19	2.33
Nitrogen		771269	777211
Oxygen		215542	221229
Carbon Dioxide		10811	1551
Methane		2368	2.48
Ethane		1.03	<0.01
Ethene		<0.01	<0.01
Propane		1.34	<0.01
Propene		<0.01	<0.01
iso-Butane		0.22	<0.01
n-Butane		0.55	<0.01
iso-Pentane		0.22	<0.01
n-Pentane		0.21	<0.01
C ₆ +		0.26	0.09
<hr/>			
C1 Index (C1/ΣC2+)		753.7	N/A
C2 Index (C2/ΣC3+)		0.49	N/A
C3 Index (C3/ΣC4+)		1.75	N/A
C4 Index (C4/C5)		2.58	N/A
ΣC2+		3.14	N/A
ATM Ratio (N2/O2)		3.58	3.51
Vol % CO2 of TG		1.08	0.16
Vol % Lt. Alk. of TG		0.24	0.00
Vol % Lt. Alk. CH4		99.8	100.00
Vol % Lt. Alk. C2+		0.15	0.00
Vol % C2+ of TG		0.00	0.00
<hr/>			
Stable Carbon Isotope Compositions (‰ VPDB)			
δ13C CH4		-50.39	nm
δ13C C2H6		nm	nm
δ13C C2H4		nm	nm
δ13C C3H8		nm	nm
δ13C C3H6		nm	nm
δ13C i-C4H10		nm	nm
δ13C n-C4H10		nm	nm
δ13C i-C5H12		nm	nm
δ13C n-C5H12		nm	nm
δ13C CO2		-11.93	nm
<hr/>			
Stable Hydrogen Isotopic Compositions (‰ VSMOW)			
δD H2		nm	nm
δD CH4		nm	nm
δD C2H6		nm	nm
δD C3H8		nm	nm
δD i-C4H10		nm	nm
δD n-C4H10		nm	nm
<hr/>			
14C Concentration (pMC)		nm	nm

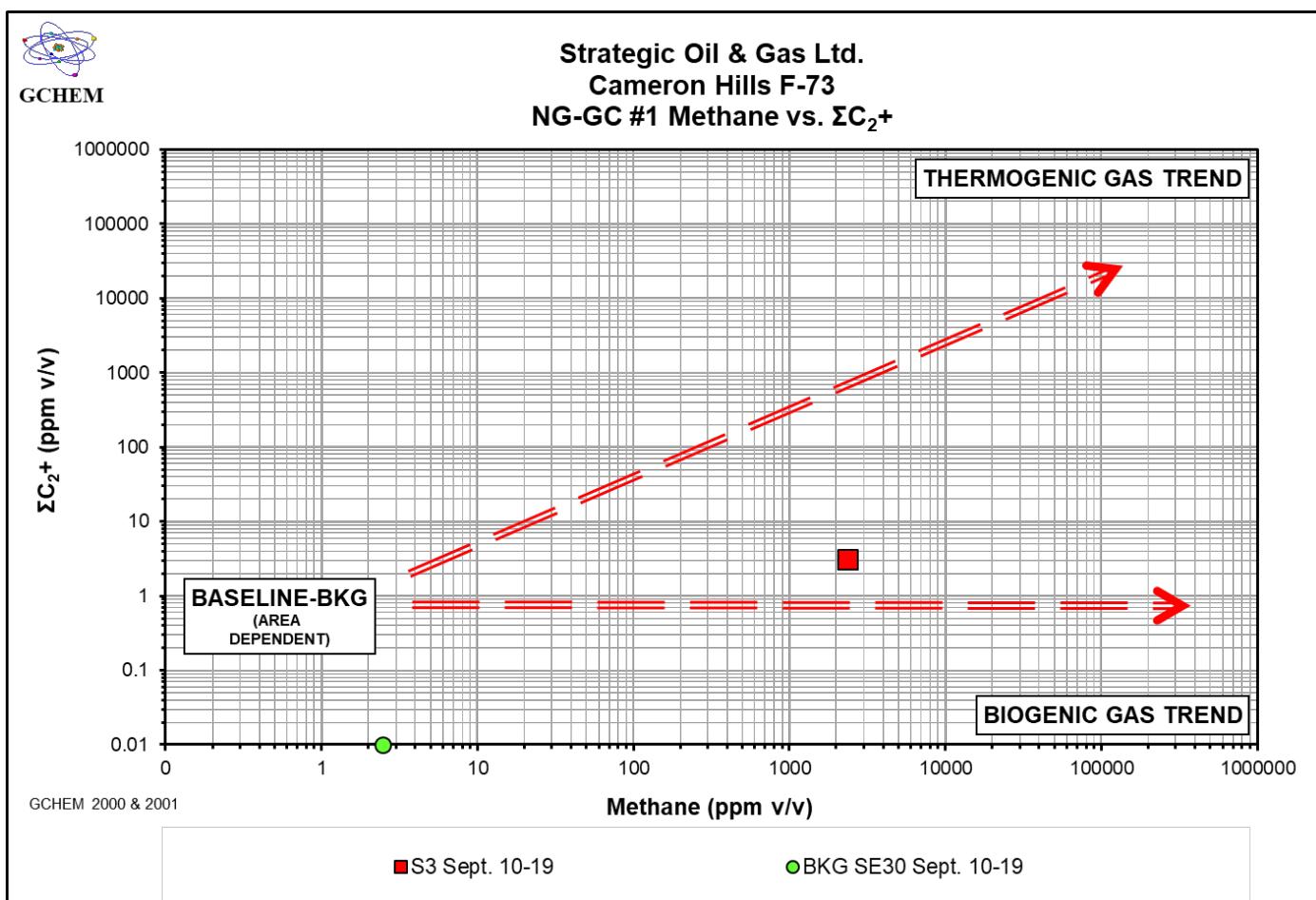


Figure-4. ΣC_2+ vs Methane. Combustible gases detected in soils and SCVs at a wellhead may result from several origins. Natural gases indicative of SCVF or AGM are thermogenic in origin (natural gas in deep reservoirs), contain high methane and C_2+ contents and plot in the Upper RH Quadrant. Low natural gas levels in background, off lease areas are naturally present in soils, vary from region to region and plot in the Lower LH Quadrant. Biogenic gases (swamp-gas) are produced by bacteria, are comprised of predominantly methane and plot in Lower RH Quadrant. Samples plotting in the Lower LH and RH do not contain SCVF or AGM and would not require down-hole remediation.

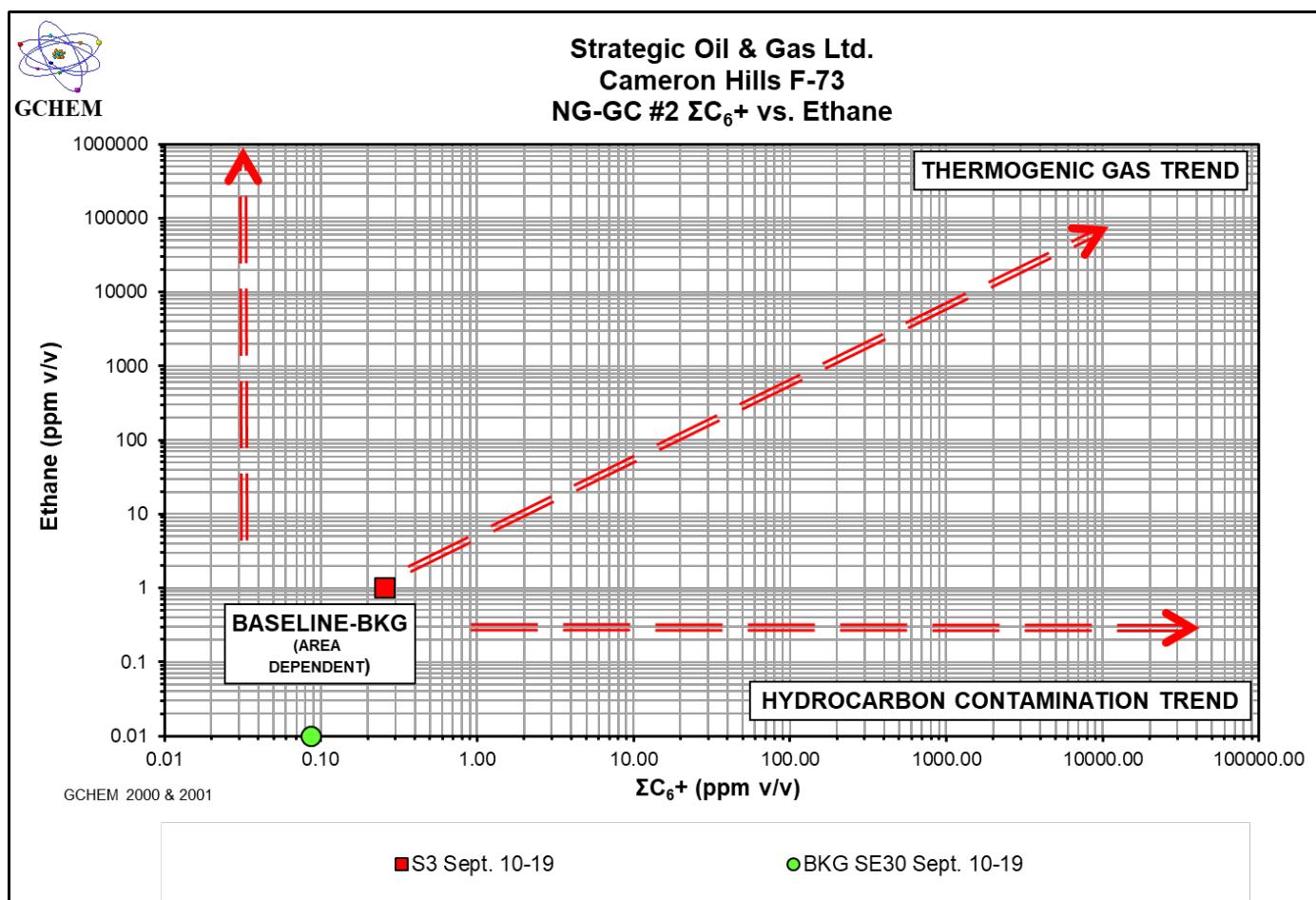


Figure 5. ΣC_6+ vs Ethane. C_6+ gases are relatively large molecules that do not readily or easily migrate in large quantities from depth upwards through subsurface fractures or micro-fractures to surface. Contamination by oil spills, fuels, and solvents is indicated by soil vapor samples that have high contents of C_6+ compounds and plot in the Lower RH Quadrant. Samples plotting in the Lower LH and RH Quadrants do not contain evidence of either SCVF or AGM and would not require downhole repair operations.

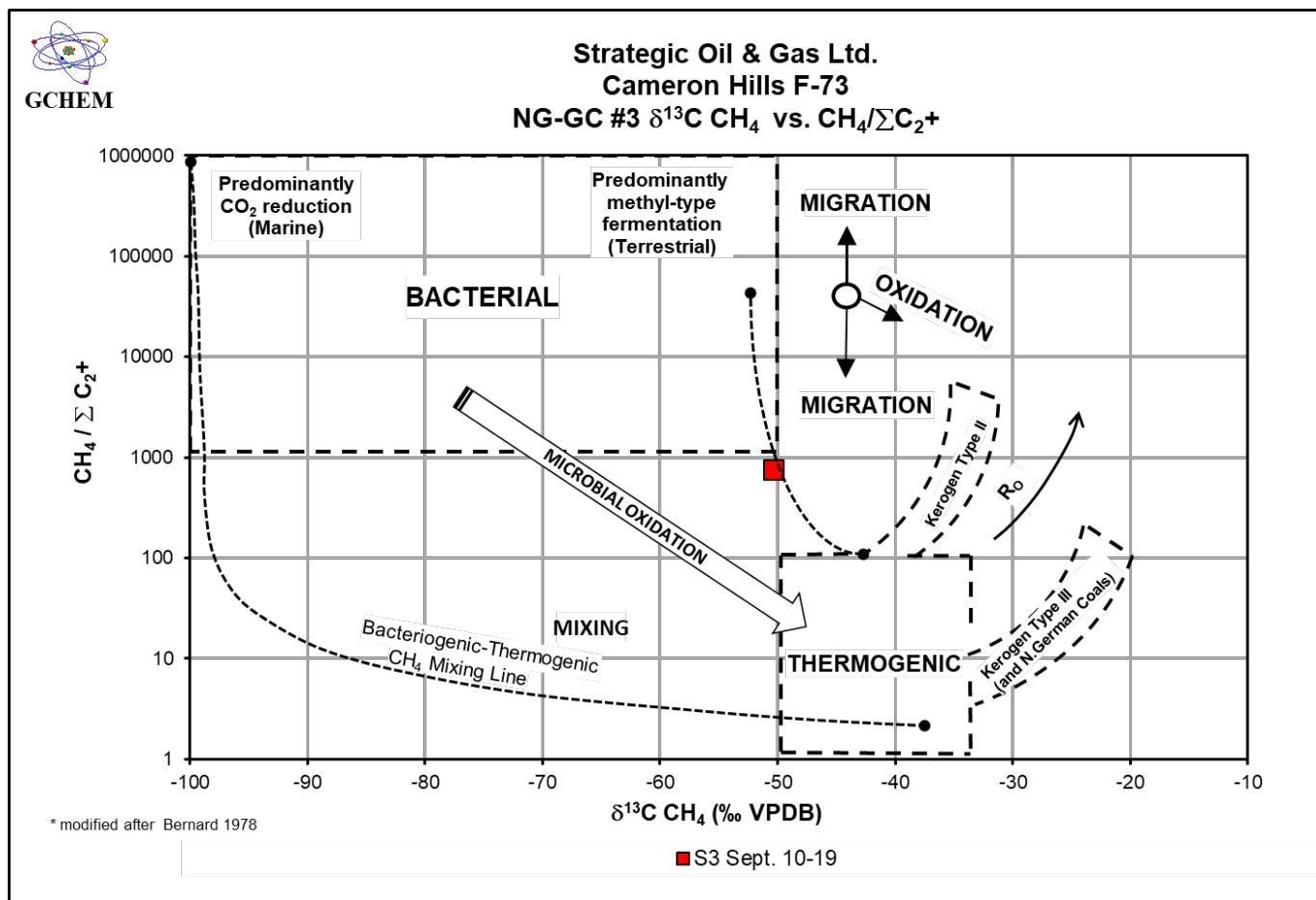


Figure 6. $\text{CH}_4 / \sum\text{C}_{2+}$ vs. $\delta^{13}\text{C CH}_4$. Thermogenic methane or methane generated by abiotic processes such as the thermal degradation of organic matter at high temperature and pressure (thermogenesis) contains enriched (less negative) $\delta^{13}\text{C}$ values ranging from -50 to -20‰ VPDB and methane relative to C_{2+} gas contents (gas wetness) less than 100. Methane gas can be generated by biotic processes such as the degradation of organic matter via CO_2 reduction or fermentation reactions generating biogenic methane. It should be noted that as a normal part of soil respiration, methane may be generated or destroyed by variable biotic pathways. Biogenic methane gas may be oxidized by bacteria resulting in an 'isotopic enriching effect' (i.e. $\delta^{13}\text{C}$ values become less negative as a result of oxidizing bacteria in soils that preferentially consume ^{12}C over ^{13}C , leaving the remaining gas enriched in ^{13}C). Since biogenic oxidization decreases the ratio between ^{12}C and ^{13}C , it may result in enriched $\delta^{13}\text{C CH}_4$ values that overlap with the MIXING or THERMOGENIC-GAS TREND. Biogenic methane may therefore contain $\delta^{13}\text{C}$ values greater than -50‰ VPDB (GCHEM Internal RD).

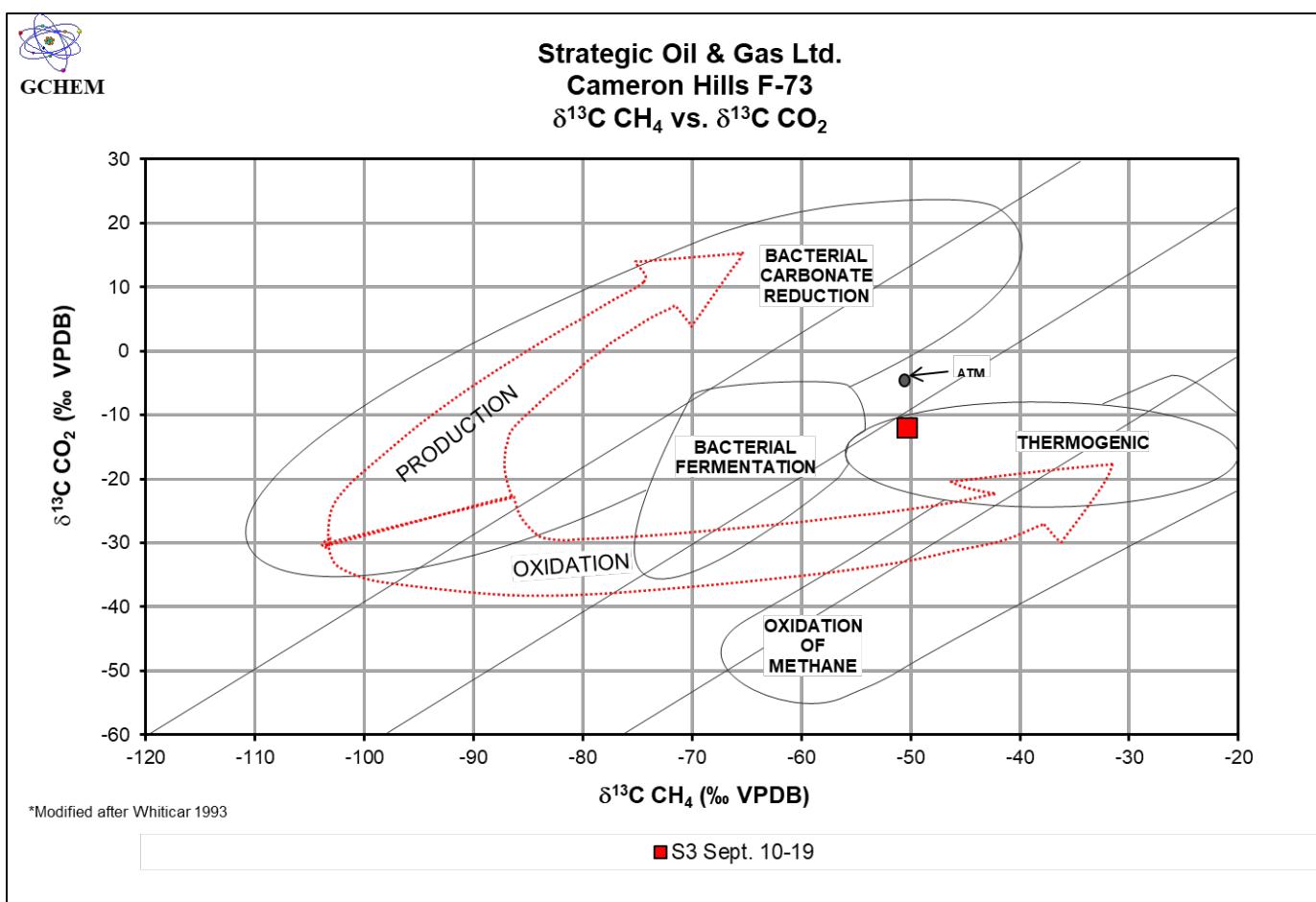


Figure 7. $\delta^{13}\text{C CO}_2$ vs. $\delta^{13}\text{C CH}_4$. Thermogenic methane or methane generated by abiotic processes such as the degradation of organic matter at high temperature and pressure contains enriched (less negative) $\delta^{13}\text{C}$ values ranging from -55 to -20‰ VPDB (or higher) and $\delta^{13}\text{C CO}_2$ values in the range of -25 to 4‰ VPDB. Methane gas may be generated by biotic processes such as the degradation of organic matter via CO_2 reduction or fermentation reactions generating biogenic methane. Biogenic methane may contain $\delta^{13}\text{C}$ values greater than -40‰ VPDB due to biogenic oxidation processes (GCHEM, in prep).

6.0 Conclusions

Soils outside casing at SOG Cameron Hills F-73 are wet and provides challenges for AGM vapor intrusion assessments. 9 of the 10 intrusive soil sites outside casing tested for combustible gas contents contained elevated methane levels that ranged from 11 to 26,850 ppm v/v. H₂S was not detected (< 1.0 ppm v/v) at any of the soil test hole sites. Flux tests conducted at N1m-E4m and in background at BGK-SE30m indicate very low gas flow rates at surface (0.000035 and 0.000033 m³/day, respectively). SOG selected sites S3m (26,850 ppm v/v) for high resolution chemical and stable carbon isotope measurements to classify combustible gas contents. Light hydrocarbon gases were dominated by methane gas while associated C₂+ gases were low and similar to background levels measured at test site BKG SE30m. Sufficient levels of CH₄ and CO₂ were available for $\delta^{13}\text{C}$ at site S3m. $\delta^{13}\text{C}$ CH₄ and $\delta^{13}\text{C}$ CO₂ at SVP S3m was -50.39 and -11.93‰ VPDB respectively.

With information available to date, SVP soil test sites S3m would be classified as 'biogenic-baseline' where CH₄ gas is the result of natural soil respirations processes via CO₂ reduction or fermentation processes generating biogenic CH₄ that is in turn further altered by secondary bacterial oxidation processes. C₂+ gases in soils near the well are low, similar to background levels and the result of natural movement of thermogenic natural gas, from reservoirs at depth, upward through fractures and micro-fractures to surface. This is a naturally occurring process prevalent in every hydrocarbon sedimentary basin in the world.



Attachment-1

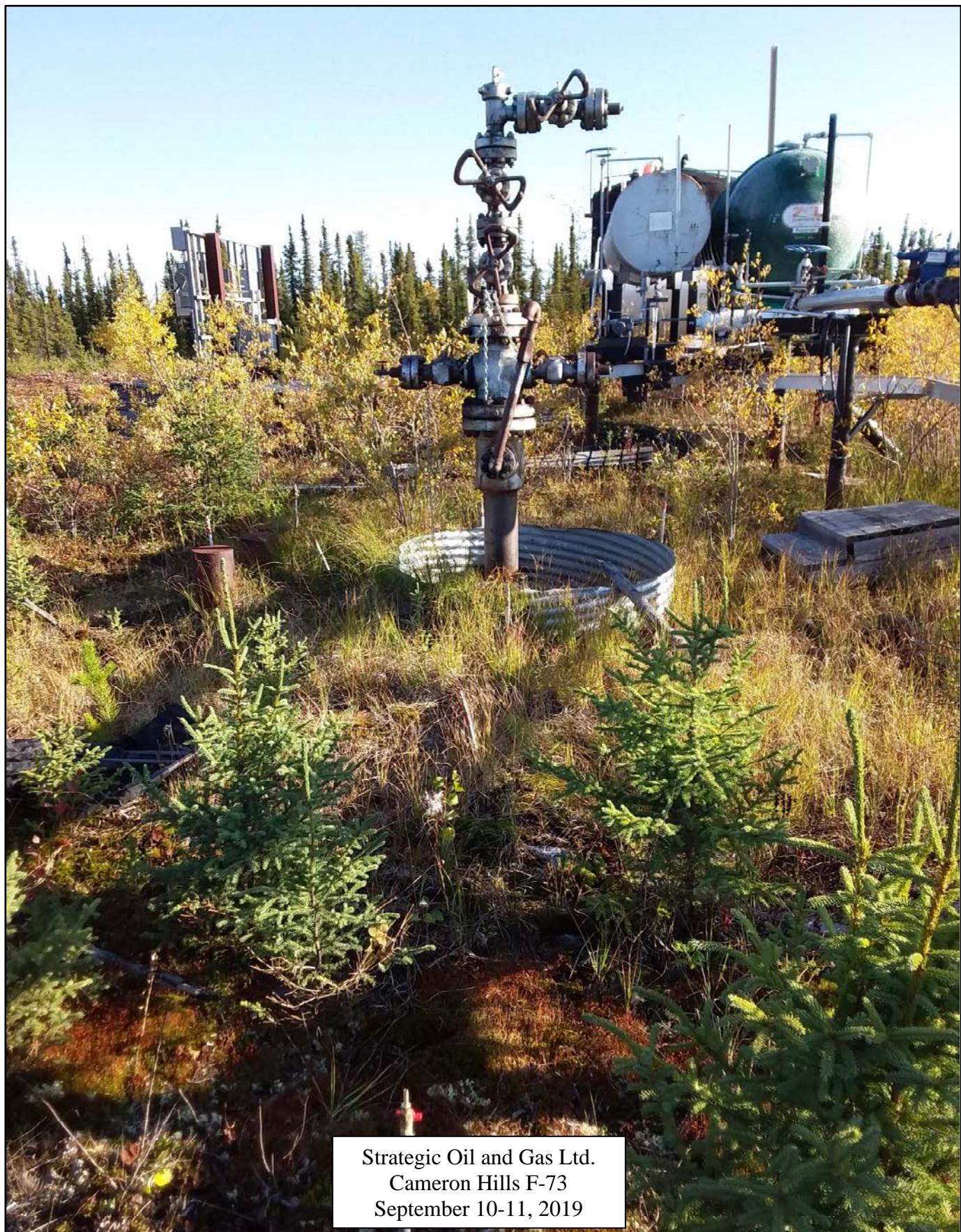
Strategic Oil & Gas Ltd.

Strategic Cameron F-73

Well Site Photographs









Attachment-2

Strategic Oil & Gas Ltd.

Strategic Cameron F-73

Chain of Custody (COC)



ENERGY FORENSICS

CHAIN OF CUSTODY

Sample Submission Form

Bay#1, 4810-62nd Ave, Lloydminster, Alberta T9V 2E9
info@gchem.ca
Tel (780) 871-4668
Fax (780) 809-4633

GCHEM LTD.

GCHEM Ltd. Project# _____

Client Information

Company Strategic
Address _____
City/Prov. _____
Postal Code _____
Client Contact _____
Phone # _____
Fax # _____
E-Mail _____

Billing/Report Information

Company _____
Address _____
City/Prov. _____
Postal Code _____
Client Contact _____
Phone # _____
Fax # _____
E-mail _____

Services Needed (TAT)

*Standard 5-7 Days
**Rush 48hrs.
***Priority Rush 24hrs.

(*) Working Days
(**) Call for Pricing and Advance Notice

Analysis To Be Performed	High Resolution Compositional H ₂ S Analysis
IRMS δ ₁₃ C Analysis	Produced Water Forensic Suite
IRMS δ ₂ H (D) Analysis	
High Resolution Compositional Analysis	

No.	Location	Sample Identifier	Sample Time	Date Sampled	Pressure Received	Actual Pressure	Container Type	Qty.	Sample Volume	Media Type
1	E73	Soil Gas - Initial	2019	Sep 10/19	1000	900	Glass	4		
2		Background - Initial	2019	Sep 10/19	1000	900	Glass	3		
3		Soil gas - Chamber	2019	Sep 10/19	1000	900	Glass	1		
4		Soil gas	2019	Sep 10/19	1000	900	Glass	1		
5		Background - Chamber	2019	Sep 10/19	1000	900	Glass	1		
6		Soil gas - Parallel/Control	2019	Sep 11/19	1000	900	Glass	5		
7		Background - Final	2019	Sep 11/19	1000	900	Glass	1		
8										
9										
10										

Sampled By Brian

A/F/PO # _____

Comments

Relinquished By: Walt Date/Time: Sep 16, 2019

Date/Time:

Relinquished To:

Date/Time:

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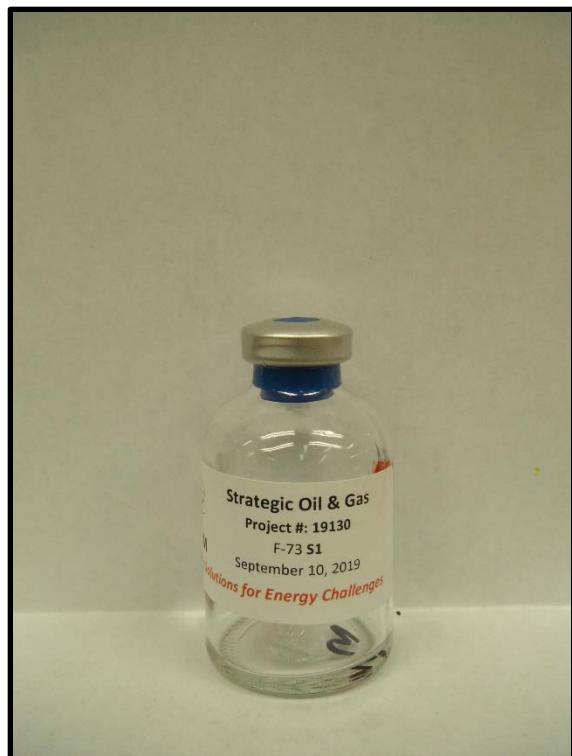
Attachment-3

Strategic Oil & Gas Ltd.

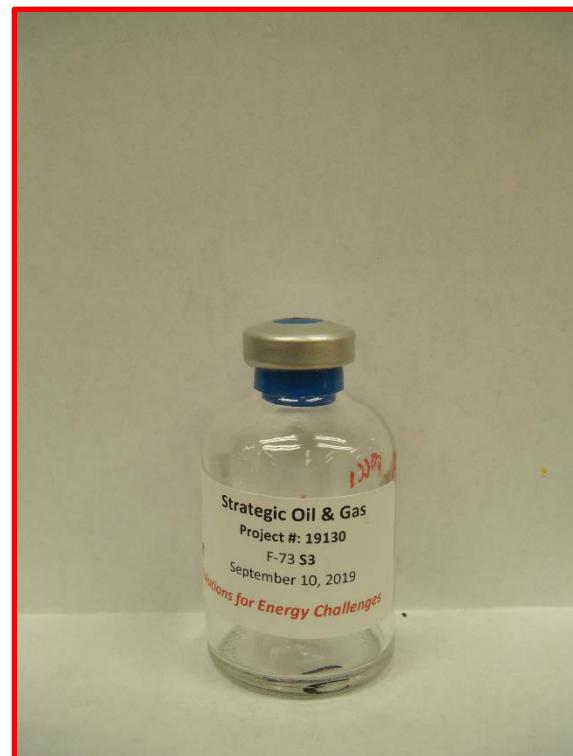
Strategic Cameron F-73

Gas Sample Containers

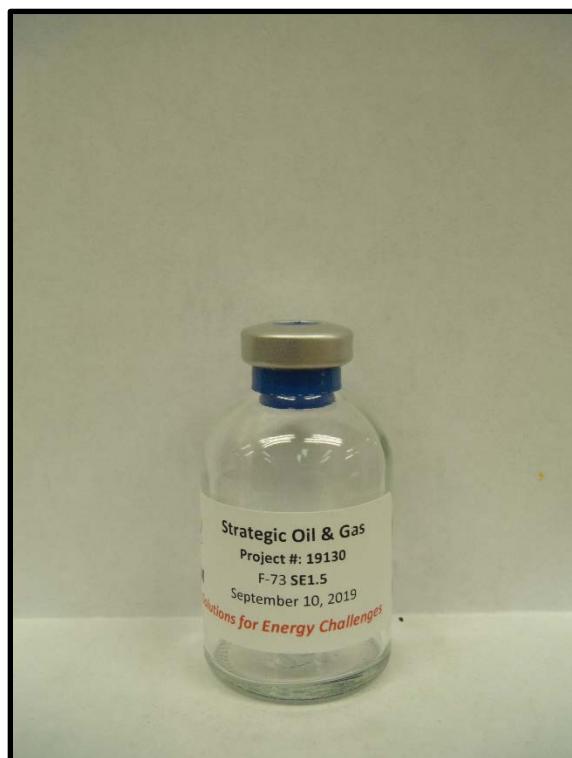
Photographs



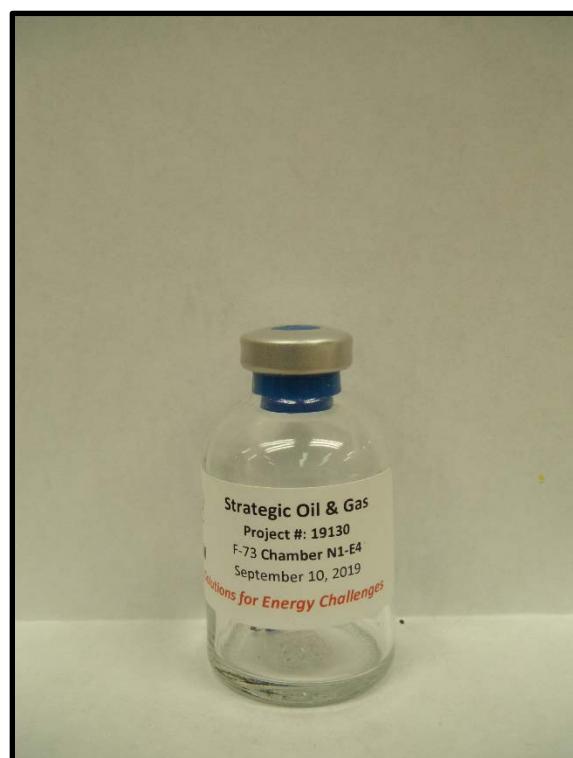
S 1m
September 10, 2019



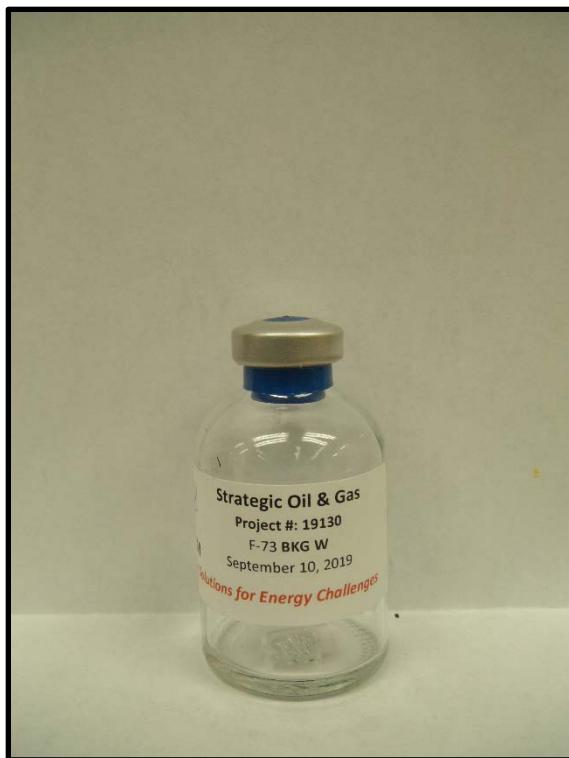
S 3m
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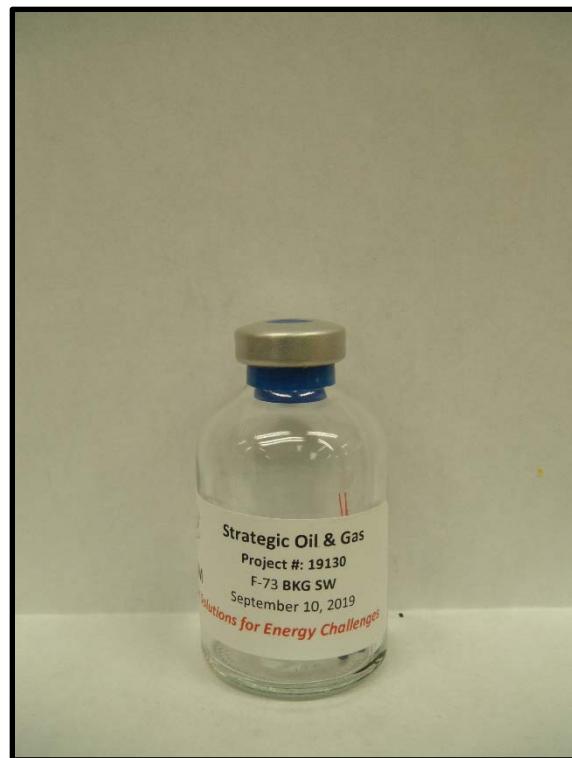
SE 1.5m
September 10, 2019



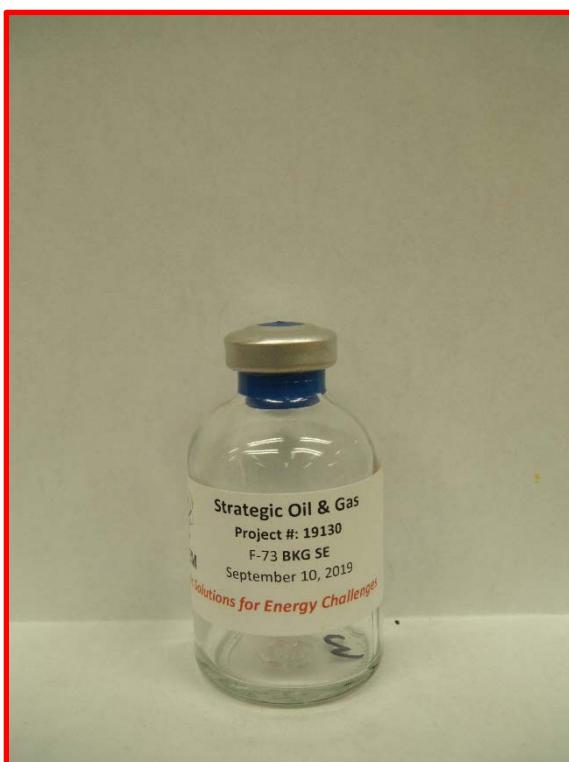
N1m-E4m Chamber
September 10, 2019



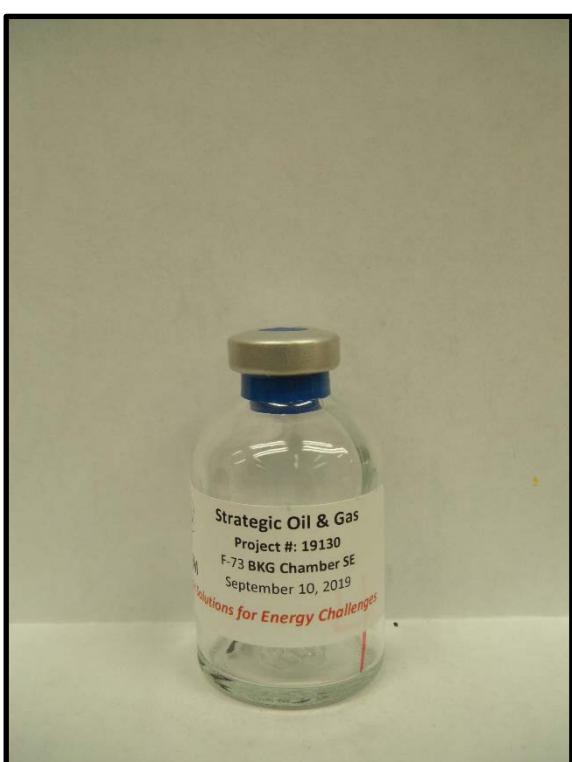
BKG W30m
September 10, 2019



BKG SW30m
September 10, 2019



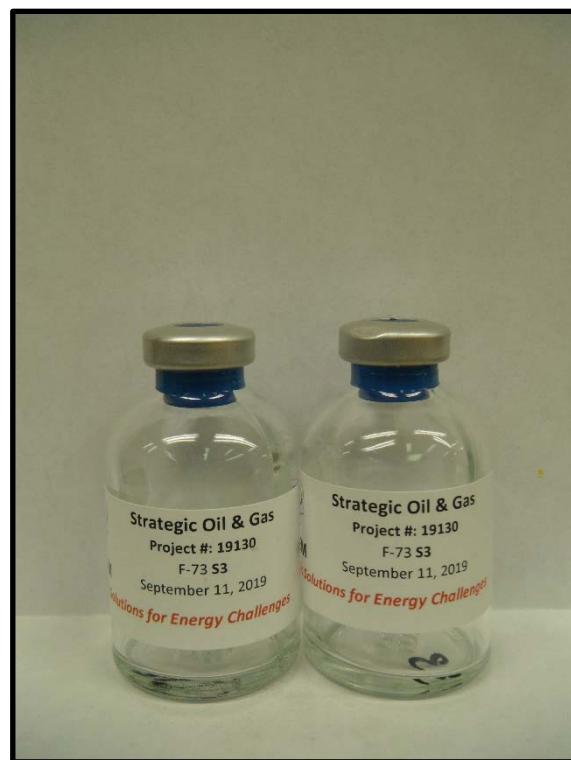
BKG SE30m
September 10, 2019



BKG Chamber SE
September 10, 2019



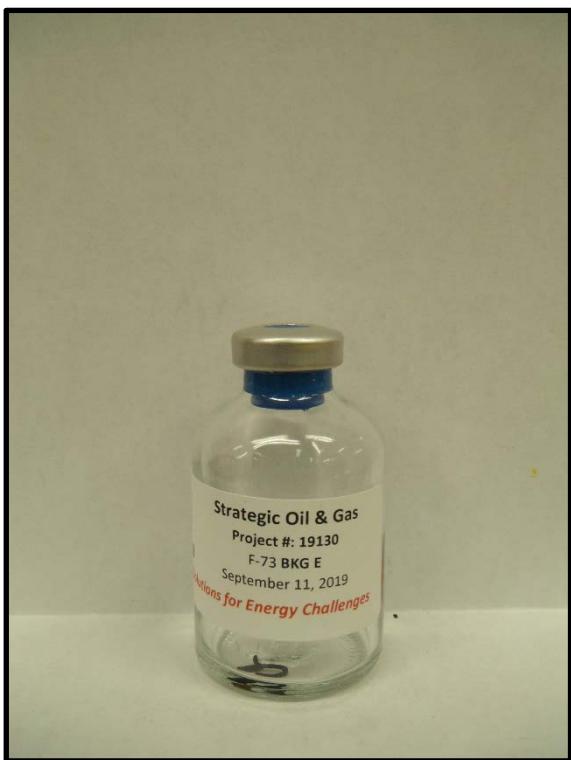
S 1m
September 11, 2019



S 3m
September 11, 2019



SE 1.5m
September 11, 2019



BKG E
September 11, 2019

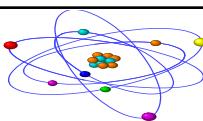


Attachment-4

Strategic Oil & Gas Ltd.

Strategic Cameron F-73

Gas Analysis Data Sheets (GADS)



GCHEM LTD.

HIGH RESOLUTION GAS ANALYSIS CARBON ISOTOPE ANALYSIS HYDROGEN ISOTOPE ANALYSIS

Sampling Company	GCHEM Ltd.	Lab Sample No.	19130-08
Date Tested	September 10, 2019	Test Type	Soil gas
Operator Name	Strategic Oil & Gas	Sample Container Type	Glass Bottle
Unique Well Identifier	F-73	Sampling Point	S3
Well Name	not provided	Test Intervals or Perfs mKB	N/A
Field or Area	not provided	Date Received	September 16, 2019
Pool or Zone	not provided	Date Reported	October 9, 2019
Well License	not provided	Entered By	Xiaolong Wang
H2S Level (Observed at Site)	not provided	Reviewed By	Brad Johnston

Sample Handling Conditions

	Source/Sampled	Received
Pressure (kPa)	N/A	52
Temperature (°C)	N/A	20

Other Information:

Laboratory Analysis

Component	HRGC Analysis As Received Mol Frac.	Air Free As received Mol Frac.	Air Free / Acid Free As Received Mol Frac.	Carbon Isotope Analysis ‰ VPDB	Hydrogen Isotope Analysis ‰ VSMOW	HRGC Analysis As Received ppm v/v
Neon	0.000020	0.001520	0.001520			20.05
Hydrogen	0.000003	0.000258	0.000258			3.40
Helium	0.000002	0.000166	0.000166			2.19
Nitrogen	0.771269	0.000000	0.000000			771269
Oxygen	0.215542	0.000001	0.000001			215542
Carbon Dioxide	0.010811	0.819715	0.819715	-11.93		10811
Carbonyl Sulphide	nm	nm	nm			nm
Hydrogen Sulphide	nm	nm	nm			nm
Methyl Mercaptan	nm	nm	nm			nm
Ethyl Mercaptan	nm	nm	nm			nm
Thiophene	nm	nm	nm			nm
Dimethyl Disulphide	nm	nm	nm			nm
Methane	0.002368	0.179569	0.179569	-50.39		2368
Ethane	0.000001	0.000078	0.000078			1.03
Ethene	0.000000	0.000000	0.000000			<0.01
Propane	0.000001	0.000102	0.000102			1.34
Propene	0.000000	0.000000	0.000000			<0.01
iso-Butane	0.000000	0.000017	0.000017			0.22
n-Butane	0.000001	0.000042	0.000042			0.55
iso-Pentane	0.000000	0.000017	0.000017			0.22
n-Pentane	0.000000	0.000016	0.000016			0.21
C ₆ +	0.000000	0.000020	0.000020			0.26
TOTAL	1.000000	1.000000	1.000000			1000000

Properties

Compositional Indicies		Real Gross Heating Value (mj/m3) @15°C and 101.35 kPa		Relative Density	
Vol % Hydrocarbons	0.24	Air Free	Moisture and Acid Gas Free	Calc. Mol. Mass Ratio	Calc. Relative Density
Vol % CH ₄	99.85	as received		1.0019	1.0019
Vol % C ₂ +	0.00				
CH ₄ / \sum C ₂ +	753.7	0.09	6.84		
C ₂ / \sum C ₃ +	0.49				
C ₃ / \sum n-C ₄ +	1.75				

Pseudo Critical Properties

	As Received	Acid Gas Free
pPc (kPa)	3792	6879
pTc (°K)	134	284

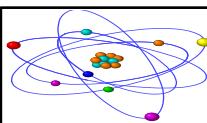
Geological Origin of Natural Gas

Geological Formation	Depth Range (MD from KB of Well)	Probable Depth (MD from KB of Well)

Comments

Forensic Solutions for Oilfield Challenges
GCHEM Ltd. Bay #1, 4810-62 Avenue Lloydminster, AB T9V 2E9 Tel: (780) 871-4668 Fax: (780) 808-8883 e-mail: info@gchem.ca www.gchem.ca GPA 2145-09. Revision 1.3, August 1, 2016





GCHEM LTD.

HIGH RESOLUTION GAS ANALYSIS CARBON ISOTOPE ANALYSIS HYDROGEN ISOTOPE ANALYSIS

Sampling Company	GCHEM Ltd.	Lab Sample No.	19130-04
Date Tested	September 10, 2019	Test Type	Soil gas
Operator Name	Strategic Oil & Gas	Sample Container Type	Glass Bottle
Unique Well Identifier	F-73	Sampling Point	BKG SE
Well Name	not provided	Test Intervals or Perfs mKB	N/A
Field or Area	not provided	Date Received	September 16, 2019
Pool or Zone	not provided	Date Reported	October 9, 2019
Well License	not provided	Entered By	Xiaolong Wang
H2S Level (Observed at Site)	not provided	Reviewed By	Brad Johnston

Sample Handling Conditions

	Source/Sampled	Received
Pressure (kPa)	N/A	62
Temperature (°C)	N/A	20

Other Information:

Laboratory Analysis

Component	HRGC Analysis As Received Mol Frac.	Air Free As received Mol Frac.	Air Free / Acid Free As Received Mol Frac.	Carbon Isotope Analysis ‰ VPDB	Hydrogen Isotope Analysis ‰ VSMOW	HRGC Analysis As Received ppm v/v
Neon	0.000020	0.013100	0.013100			20.43
Hydrogen	0.000003	0.002006	0.002006			3.13
Helium	0.000002	0.001496	0.001496			2.33
Nitrogen	0.777211	0.000000	0.000000			777211
Oxygen	0.221229	0.000009	0.000009			221229
Carbon Dioxide	0.001551	0.994845	0.994845			1551
Carbonyl Sulphide	nm	nm	nm			nm
Hydrogen Sulphide	nm	nm	nm			nm
Methyl Mercaptan	nm	nm	nm			nm
Ethyl Mercaptan	nm	nm	nm			nm
Thiophene	nm	nm	nm			nm
Dimethyl Disulphide	nm	nm	nm			nm
Methane	0.000002	0.001590	0.001590			2.48
Ethane	0.000000	0.000000	0.000000			<0.01
Ethene	0.000000	0.000000	0.000000			<0.01
Propane	0.000000	0.000000	0.000000			<0.01
Propene	0.000000	0.000000	0.000000			<0.01
iso-Butane	0.000000	0.000000	0.000000			<0.01
n-Butane	0.000000	0.000000	0.000000			<0.01
iso-Pentane	0.000000	0.000000	0.000000			<0.01
n-Pentane	0.000000	0.000000	0.000000			<0.01
C ₆ +	0.000000	0.000056	0.000056			0.09
TOTAL	1.000000	1.000000	1.000000			1000000

Properties

Compositional Indices		Real Gross Heating Value (mJ/m3) @15°C and 101.35 kPa		Relative Density	
Vol % Hydrocarbons	0.00	Air Free	Moisture and	Calc. Mol. Mass Ratio	Calc. Relative Density
Vol % CH ₄	100.0	as received	Acid Gas Free	0.9985	0.9985
Vol % C ₂ +	0.00	0.00	0.09		
CH ₄ / \sum C ₂ +	N/A				
C ₂ / \sum C ₃ +	N/A				
C ₃ / \sum n-C ₄ +	N/A				

Pseudo Critical Properties		
	As Received	Acid Gas Free
pPc (kPa)	3762	7354
pTc (°K)	133	303

Geological Origin of Natural Gas

Geological Formation	Depth Range (MD from KB of Well)	Probable Depth (MD from KB of Well)

Comments

Forensic Solutions for Oilfield Challenges
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