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Hydrogeological Characterization Report

Issaquah, Washington

Prepared for

City of Issaquah, Washington 1775 – 12th Ave NW Issaquah, WA 98027

Prepared by

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1. INTRODUCTION

Geosyntec Consultants (Geosyntec) has prepared this report for the City of Issaquah (City) to document results of drilling and sampling activities completed in 2016 in the vicinity of City water supply Wells 4, 5, and 6 (Site). The water supply wells provide drinking water for the City of Issaquah.

The objectives of the field investigation were to: 1) develop an understanding of the distribution of perfluorooctanesulfonic acid (PFOS) in groundwater near City Well 4; and 2) establish a conceptual model of how PFOS entered and is transported in the upper aquifer that is pumped by City Well 4. PFOS and perfluorooctanoic acid (PFOA) are two isomers of a larger class of compounds known as perfluorinated compounds (PFCs) or perfluoroalkyl acids (PFAAs). An overview of PFAAs is provided in our earlier Phase 1 investigation report (Geosyntec, 2016a). The term PFCs will be used herein to describe this group of compounds¹.

On May 25th, 2016 the Environmental Protection Agency (EPA) released a drinking water health advisory for PFOS which lowered the advisory level for PFOS from 0.2 micrograms per liter (μ g/L) to 0.07 μ g/L, and for PFOA from 0.4 μ g/L also to 0.07 μ g/L (EPA, 2016a, b). EPA also recommended that when these two compounds occur together in a drinking water source that the sum of the PFOS and PFOA concentrations be compared to 0.07 μ g/L. In other words, the drinking water health advisory level applies to both PFOS and PFOA independently and summed together.

The investigation included:

- The installation of seven groundwater monitoring wells: MW01, MW02, MW03, MW04, MW05, MW06, and MW07 (Figure 1);
- Groundwater quality sampling at the seven new wells and several existing wells in the vicinity of the City's water supply wells;
- Completion of a video log in an existing City test well near Well 5;
- Continuous monitoring of groundwater levels in six wells;

¹ The term "PFCs" is also used to describe compounds associated with ozone depletion.



- Sampling of soil at Eastside Fire and Rescue; and
- Surface water sampling in Issaquah Creek.

A draft work plan was prepared and submitted to the City on 17 May 2016 (Work Plan) describing the well installation and groundwater monitoring activities; however, through the course of the investigation the field program was modified based on the first round of sampling data. Drilling, sampling, and monitoring procedures were generally completed as described in the Work Plan.

2. DRILLING AND MONITORING WELL INSTALLATION

Monitoring well boreholes were drilled using track-mounted Sonic[™] drilling rigs operated by Holt Services, Inc. (Edgewood, WA), and Cascade Environmental Services (Woodinville, WA). The monitoring wells were installed in two phases, MW01 through MW05 were completed in May 2016, and MW06 and MW07 were completed in October 2016. Continuous sample cores were obtained during drilling to characterize subsurface materials, conduct field screening, and identify zones for depth-discrete grab water samples. Borehole depths ranged from 80 feet below ground surface (bgs) to 110 feet bgs, as summarized in Table 1. Borehole logs, including field screening results and sample depths, are included in Appendix B.

2.1 Geology

The surficial geology in the vicinity of the City of Issaquah is shown in map form on Figure 2. The surficial materials are predominantly glacial sediments, but there is an outcrop of bedrock in the southwest portion of the City at the intersection of Sunset Way and Newport way.

Cross-sections that show our interpretation of the geology based on the borehole drilling are presented on Figures 2A and 2B. Subsurface materials encountered during this investigation generally consisted of:

• Brown sand or sandy silt to approximately 20 feet bgs, underlain by a grey to grey-green sandy silt from approximately 20 to 40 feet bgs. These two units are interpreted to be shallow alluvium (Qa) underlain by ice contact deposits (Qvi).

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- A loose, grey sand was encountered from approximately 40 feet bgs to 60 or 70 feet bgs (ice contact deposits [Qvi]). This sand unit was difficult to drill through due to heaving sand conditions. The heaving sand unit was underlain by a grey sandy silt with occasional wood fragments approximately 10 to 20 feet in thickness (60 to 70 feet bgs to 70 to 80 feet bgs). Some organic material (wood fragments) and one small section of peat were encountered near the base of the ice contact deposits.
- Brown to grey brown sand to silty sand interpreted to be advance outwash sediments (Qva) were typically encountered at 70 to 80 feet bgs up to the maximum depth explored of 100 feet bgs. At MW02, the brown gravelly sand (Qva) is thinner than the other boreholes and advance outwash sediments were not encountered at MW01, west of MW02.
- These materials correspond to geologic units in the area, which are outlined in the Work Plan (Geosyntec, 2016b), and consist of the following formations (from shallowest to deepest):
 - o Shallow alluvium (Qa);
 - o Ice Contact Deposits (Qvi);
 - o Advance Outwash (Qva);
 - Older glacial deposits (Qpff); and Olympian Sediment (Qob).

These units have an apparent northward hydrostratigraphic slope of about 1.4% (Figure 2A). Coarser grained sand and gravel generally was present to the south and finer sand and silt to the north, consistent with deposition in a deltaic (fining downslope) type setting. The advance outwash (Qva) sediments intersected in MW02, MW03, MW04, MW05, MW06, and MW07 were each consistent with geology described in other City wells. These glacial advance outwash sediments were thinner at MW02. Glacial advance outwash sediments were not observed at MW01. This suggests that the advance outwash pinches out in the vicinity of MW02.

2.2 Discrete-Depth Groundwater Sampling

Groundwater samples were obtained from the boreholes during drilling to evaluate PFC concentrations at first encountered groundwater and at additional selected depths for vertical profiling. The sample results were collected in an open borehole and were not intended to substitute for results from a constructed monitoring well.



Sampling depths and results of the discrete-depth samples are discussed in more detail in Appendix A.

2.3 Monitoring Well Installation

A monitoring well was constructed in each borehole drilled at a depth interval determined in the field. Monitoring wells are constructed of 2" polyvinyl chloride (PVC) well casing with 0.010" slotted PVC well screens. A prepack well screen and above ground monument were used at MW02 due to heaving sand and artesian (flowing) conditions.

Borehole and monitoring well construction logs are included as Appendix B. The City completed a survey of the monitoring well locations and elevations on 10 June and 12 November 2016. Monitoring well construction details and survey data are summarized in Table 1.

3. GROUNDWATER ELEVATION MONITORING

Self-contained pressure transducers (Van Essen Divers or Divers) were initially installed in five monitoring wells (MW01 through MW05) and City of Issaquah Well COI-TW-3. The transducers in MW01 and MW02 were then moved to MW06 and MW07. In addition to the Divers, a pressure transducer that records barometric pressure (Baro-Diver) was installed at MW02 on the underside of the well monument cover. The Baro-Diver is used to compensate the groundwater pressure recorded by the Divers for changes in barometric pressure.

Manual groundwater measurements in the monitoring wells were measured prior to installation of the transducers and ranged from 55 feet above mean sea level (amsl) to 65 feet amsl (Table 2). The general direction of groundwater flow is from southeast to northwest.

The Divers began recording data on 2 June 2016. Diver operation was verified against field measurements by suspending cable length and manual depth-togroundwater measurements compared to recorded groundwater pressures. Groundwater pressures recorded by the Divers were corrected for barometric pressure and converted to groundwater elevations using manual depth-to-water measurements and the surveyed top of casing elevations at each well.

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The Diver data collected for the 2 June through the 17 October 2016 are presented as a time-series in Figure 4. The Figure shows the general seasonal trend of groundwater levels between June and October 2016. Groundwater levels generally decline about 2 to 4 feet during the course of the summer and rise again in September. The large changes in groundwater level shown at MW02 and MW03 in June and July are from testing at City Well 4. Data are recorded at 15 minute intervals in the transducer, but Figure 4 presents the average daily groundwater level for clarity. There is a high frequency water level fluctuation observed within the course of a day at MW02, MW03, MW04, and MW05. The fluctuations correspond to pumping operations at Sammamish Plateau Sewer and Water District (SPSWD) Wells 7 and 8, which are completed in the same aquifer. The fluctuation is not observed in the groundwater elevation record for MW01, which is screened above the advance outwash aquifer. The fluctuation is largest in MW04, causing changes up to one foot in groundwater elevation during the course of a day.

4. GROUNDWATER QUALITY MONITORING

Groundwater sampling was completed at monitoring and production wells (Figure 1), as follows:

- Darigold well (ABY249) was sampled on 4 May 2016;
- SPSWD wells 7.3, VT-2.1, VT-2.2, VT-2.3 were sampled on 10 May 2016;
- City of Issaquah wells COI6-TMW1s (formerly COI-MW1) and COI-TW-3 were sampled on 17 May 2016; and City Well 5 was sampled on 6, 13, and 27 July 2016, and
- New City of Issaquah monitoring wells MW01, MW02, MW03, MW04, and MW05 were sampled on 7 June 2016. Monitoring wells MW03, MW04, MW05 were sampled weekly from 21 June through 27 July 2016.
- Monitoring wells MW01 through MW07 were sampled on 17 October 2016.

Groundwater monitoring was completed using EPA protocols, including low flow purging and sampling of the new wells as outlined in the Work Plan (Geosyntec, 2016b). Sample preservation and handling procedures were developed with the laboratory to prevent cross-contamination with potential PFC-containing materials (i.e. Teflon), as described in the quality assurance project plan attached to the Work Plan (Geosyntec, 2016b), and as outlined in Table 1 of the Interim Guideline on the Assessment and Management of Perfluoroalkyl and Polyfluoroalkyl Substances (State of Western Australia, 2016).

The SPSWD and City of Issaquah monitoring wells were purged and sampled using a peristaltic pump and disposable tubing. Approximately three well volumes were purged from each well prior to sampling. The Darigold well is a water supply well and was pumping during sampling, so a grab groundwater sample was obtained from a sample port at the well head after flushing for two minutes. Low-flow purging of the new groundwater monitoring wells was completed using a peristaltic pump and disposable tubing along with a Horiba water quality instrument to collect stabilization parameters. Following stabilization, groundwater samples were obtained for analytical testing of:

- PFCs using EPA Method 537;
- Cations (calcium, chloride, magnesium, potassium, and sodium) by EPA Method 200.7;
- Anions (sulfate and nitrate) by EPA Method 300;
- Alkalinity by standard method (SM) 2320B;
- pH by SM 4500-HB;
- Specific conductance by SM 2510B; and
- Benzene, toluene, ethylbenzene, total xylenes (BTEX), and methyl tertbutyl ether (MTBE) by EPA Method 524.2.

Prior to sampling, depth to groundwater was measured using an electronic depth to water meter.

Analytical laboratory reports are included in Appendix C along with Data Quality Review Memos. Results of the data quality review indicate that the analytical data are acceptable for their intended use.

In addition, SPSWD Wells 7, 8, 9 were sampled for PFCs by SPSWD on 15 and 20 June, and 26 July 2016 and analyzed by Eurofins Laboratory in Monrovia CA. These results are included in Table 3 and on Figure 5.

5. GROUNDWATER SAMPLING ANALYTICAL RESULTS

Analytical results for BTEX and cations/anions results are summarized in Table 3, and PFC results for groundwater are summarized in Table 4.

5.1 <u>Perfluorinated Compounds (PFCs) Results</u>

PFCs, including PFOS, have been detected in thirteen wells (Table 2) in the Lower Issaquah Valley, including City Wells 4 and 5, SPSWD Wells 7, 8, and 7.3, COI6-TMW1s (COI-MW1), the Darigold Well, MW01, MW03, MW04, MW05, MW06, and MW07. PFOS Results are summarized as follows:

- PFOS was detected above the EPA advisory level of 0.07 μg/L in MW03 (0.46 μg/L), MW05 (0.39 μg/L), MW06 (2.2 μg/L), and City Well 4.
- PFOS was detected below the EPA advisory level at COI-MW1, Darigold, City Well 5, and MW07.
- PFOS concentrations below the EPA advisory level were detected by SPSWD in Well 7 (0.019J and 0.015 µg/L) and Well 8 (0.029) µg/L.
- PFOS was not detected in MW01, MW02, and MW04.
- PFOA was detected at concentrations above the EPA advisory level at MW06 (0.08 μ g/L). PFOA was detected at concentrations below the EPA advisory level (between 0.003 to 0.03 μ g/L) at MW01, MW03, MW04, MW05, and MW07.

Additional PFCs (i.e. other isomers of the larger class of compounds) were detected in City Wells 4 and 5, MW03, MW05, and MW06. These other perfluorinated compounds do not have associated EPA advisory levels.

The distribution and sampling history of PFOS in the various monitoring wells is shown in a map format on Figure 5. Note, this figure shows both PFOS and PFOA (where detected).

5.2 Anions/Cations Ratios

The distribution of anions/cations at MW03, MW04, MW05, and MW06 are similar to each other indicating similar groundwater chemistry (i.e. same aquifer or recharge source). Slightly different cation compositions are observed at MW01, MW02, and

MW07. Water quality parameters are plotted in Figures 6 and 7 to evaluate their relative concentrations.

6. OBSERVATION WELL 5 VIDEO SURVEY

PFOS is present in City Well 5 at a concentration of 0.03 μ g/L. Well 5 is completed in a deeper aquifer at 240 to 412 feet bgs. The deeper aquifer is separated from the advance outwash aquifer by approximately 100 feet of silt.

A video survey of an older observation well adjacent to City well 5 (Well 5OBS) was completed to assess the integrity of the well and determine whether it could be a conduit for PFCs to move from the shallow aquifer to the deep aquifer. Well 5OBS was drilled in the early 1980s as a test well for the City's deep production well. Well 5OBS was not completed with a well screen, but has a slotted casing from approximately 330 to 450 feet below ground surface.

The video survey of well 5OBS did not show signs of leakage, cracks, or breaks in the well casing. A cement plug was encountered at approximately 307 feet bgs that prevented the video camera from advancing into the completion interval. The purpose, extent, and date of installation of the plug are unknown. A photo-log showing some of the joints and welds in well 5OBS is provided in Appendix D.

7. SOIL SAMPLING

Based on the initial groundwater sampling results, we interpreted a south-trending distribution of PFOS in groundwater from City Well 4, toward MW05. Based on groundwater flow directions, a potential source location included the Eastside Fire and Rescue (EF&R) property, located at 175 Newport Way NW. This location site appears to be along a groundwater flow path that intersects both MW05 and City Well 4, and is located within the 10-year time of travel delineated in the City's Wellhead Protection Plan.

The City of Issaquah initially contacted EF&R in July and requested permission to collect soil samples, which was granted. On 22 July 2016, a Geosyntec technician collected three soil samples at the site. On 27 July 2016, Geosyntec and the City of Issaquah met with EF&R to discuss groundwater results in the monitoring wells. On 05 August 2016, Geosyntec and the City of Issaquah met with EF&R again to discuss historical fire training and fire response activity.

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The purpose of the sampling at 175 Newport Way NW to determine the potential presence or absence of PFCs in soil at the property. The sampling effort was not intended to characterize the history, magnitude, distribution, or extent of PFCs on the site.

Three soil samples were obtained on 22 July 2016 from three locations (STTA01, STTA02, and STSP01) in a gravel area on the western portion of the property (Figure 8). The soil samples were collected using hand methods (shovel and hand-auger) at depths ranging from 27 to 46.5 inches bgs. The shovel and hand auger were decontaminated between sample locations. Subsurface materials consisted of a surface gravel layer from approximately 7 to 14 inches thick, a dark grey silt at STTA01 and STTA02, and a light brown sandy silt at location STSP01. Soil samples were placed directly into laboratory supplied containers, placed into a cooler with ice and shipped to Eurofins Eaton Analytical Labs in Lancaster PA for testing of PFCs by EPA Method 537. On 19 August 2016, Geosyntec received soil sample analytical results and initiated internal data validation.

Soil sample results for 175 Newport Way NW are summarized on Table 5. PFCs, including PFOS, were detected in all three soil samples, as follows.

- PFOS was detected in all three soil samples, and concentrations ranged from 23 to 1,300 nanograms per gram (or 0.023 to 1.3 mg/kg).
- PFOA was detected in two of the three soil samples at concentrations ranging from 5 to 11 ng/g (or 0.005 to 0.011 mg/kg). These two samples were from the area at the east end of the property.

8. SURFACE WATER SAMPLING

Two surface water samples were obtained from Issaquah Creek on 27 September 2016. Samples were collected at locations upstream of the fish hatchery (SW-A) and downstream of City Wells 4 and 5 (SW-B). The sample locations are shown on Figure 9. The samples were submitted for analytical testing of PFCs by EPA Method 537.

PFCs were not detected in the surface water samples. Analytical results are summarized in Table 4.



9. SUMMARY AND CONCLUSIONS

The sub-surface geology encountered in the boreholes consists of interbedded recent alluvial deposits underlain by and interbedded with glacial sediments that generally dip northward. The surficial geology in the area also consists of predominantly glacial sediments, but there is an outcrop of bedrock at the intersection of Sunset Way and Newport way. The glacial sediments terminate against this bedrock and represent the western edge of the aquifer system in this part of the valley. The bedrock contact extends northward along Newport way and appears as a "hookshaped" feature on the map that wraps around the western portion of the property at 175 Newport Way NW.

Groundwater is typically found 20 to 30 feet below ground surface and groundwater flow is to the north, down valley towards Lake Sammamish. Silt interbeds are present in the upper 80 feet of the glacial sediments, but do not appear to be sufficiently thick nor continuous across the area to form aquitards that would limit vertical groundwater flow or connectivity through the upper aquifer.

PFOS is present in groundwater above the EPA advisory limit in City Well 4 and monitoring wells MW03, MW05, and MW06 (Figure 5). PFOS is not present at monitoring wells MW01, MW02, and MW04, and is present just above the detection limit at MW07. PFOS concentrations increase significantly in an upgradient direction from MW03 to MW06. The highest concentrations detected in this study were present in MW06 at 2.2 μ g/L (a factor of 30 times higher than the EPA advisory limit).

The extent of PFOS at concentrations greater than 0.1 μ g/L in the shallow aquifer appears to be a narrow and north-south oriented plume, as defined by detections in monitoring wells MW03, MW05, and MW06, and the absence of PFCs in MW01, MW02, MW04, and MW07. The north-south trending plume aligns parallel and west of Issaquah Creek. As shown on Figure 10, PFOS concentrations increase markedly between MW05 and MW06, which is located about 500 feet downgradient of 175 Newport Way NW, where PFOS was detected in soil. Historic use of firefighting foams at the 175 Newport Way NW, the presence of PFOS in soil, and the high PFOS detection at MW06 located just downgradient of this property, indicates this property is a source of PFCs to groundwater that should be investigated further.

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The property at 175 Newport Way NW is also located north of an eastward jutting outcrop of bedrock (a "hook-shaped" feature on the map that wraps around the western portion of the property). The configuration of the bedrock may help to concentrate infiltration and focus the flow of infiltration on the property toward MW06. Although there was a trace detection of PFOS at MW07, located upgradient of the property, it is highly unlikely that a PFOS plume capable of producing a high concentration at MW06 could exist between MW07 and the bedrock outcrop. As such, we conclude that the 175 Newport Way NW property is the primary source of PFOS contributing the plume that intersects the City's Well 4.

The concentration of a contaminant at any given point within a contaminant plume varies over time and can increase or decrease because of contaminant transport processes, the history at the source area, and the concentration of contamination at the source area. It is likely that the source at 175 Newport Way NW has reached a steady state condition, and, if no source removal actions are undertaken, concentrations at MW06 would remain high until the source area was diluted of its residual PFOS. The high concentrations of PFOS at MW06, relative to the downgradient monitoring wells, suggest that PFOS concentrations will likely increase at the City's Well 4 before they decrease. Additional modeling of contaminant transport would be necessary to quantitatively predict future PFOS concentrations at City Well 4, but they are unlikely to reach the high levels observed at MW06.

Trace levels of PFOS were detected at SP7-3, SP-7, SP-8 wells, and City Well 5 (completed in the deep aquifer), indicating that deeper transport pathways are present. The specific transport pathways for PFOS to these locations are not known and this investigation did not characterize the deeper aquifer nor potential groundwater flowpaths to these wells.

Use of fire-fighting foams at the 2002 I-90 Tanker Fire west of City Well 4, originally suspected as a source of PFOS, was eliminated as a PFOS source to City Well 4. PFOS was not detected at MW01, located directly adjacent to the fire and staging area for fire response. The water quality data and water-level responses are also different at MW01 compared to other wells. The presence of PFOS at other wells upgradient of City Well 4 also eliminates this event as a source.



10. RECOMMENDATIONS

Based on these results, Geosyntec offers the following recommendations:

- 1. Conduct a more detailed soil and groundwater investigation at the 175 Newport Way NW property to determine the extent and distribution of PFCs on that property. The investigation should include sampling of current firefighting foams on site and, if possible, any material flushed from fire-fighting equipment and stored on-site.
- Continue monitoring the recently installed monitoring wells (MW01-MW07) for water-levels and PFCs to evaluate seasonal trends in water levels and water quality.
- 3. The City's treatment system at Well 4 is currently functioning well, but we recommend that City evaluate operating procedures and determine whether adjustments to the current pumping schedule at Well 4 are warranted. This analysis should be coordinated with SPSWD so that an overall pumping strategy for the aquifer system is established. This would entail groundwater flow modeling that could also be used to evaluate contaminant transport. A simple well interference analysis that focuses solely on well pumping rates and hydraulic gradients could also be sufficient.
- 4. Once the 175 Newport Way NW source area has been better characterized and potential approaches to source removal have been evaluated, groundwater transport modeling may be useful to evaluate the longer term fate and transport of PFOS into the aquifer system.



11. REFERENCES

Geosyntec Consultants, 2016a. Phase 1 Hydrogeological Characterization. Issaquah, Washington, 4 March 2016.

Geosyntec Consultants, 2016b. Draft Hydrogeological Characterization Work Plan. Issaquah, Washington.

State of Western Australia, 2016. Interim Guideline on the Assessment and Management of Perfluoroalkyl and Polyfluoroalkyl Substances (PFAS) Contaminated Sites Guidelines, February 2016.

U.S. Environmental Protection Agency, 2016a. Drinking Water Health Advisory for Perfluorooctane Sulfonate (PFOS), EPA Document Number: 822-R-16-004, May 2016.

U.S. Environmental Protection Agency, 2016b. Drinking Water Health Advisory for Perfluorooctanoic Acid (PFOA), EPA Document Number: 822-R-16-005, May 2016.

TABLES

Table 1Monitoring Well ConstructionCity of Issaquah Hydrogeologic Assessment

		Well	Boring	Completed	Ground	Top of Casing	Top of Screen	Bottom of Screen	Top of Screen	Bottom of Screen
Well	Date	Diameter	Depth	Depth	Elevation	Elevation	Depth	Depth	Elevation	Elevation
Name	Installed	(in)	(ft)	(ft)	(ft amsl)	(ft amsl)	(ft bgs)	(ft bgs)	(ft amsl)	(ft amsl)
MW01	5/26/2016	2	100	38	58.4	58.4	28	38	30.4	20.4
MW02	6/2/2016	2	100	90	59.7	62.8	70	90	-10.3	-30.3
MW03	5/24/2016	2	100	98	63.2	62.9	78	98	-14.8	-34.8
MW04	5/27/2016	2	90	90	73.3	73.1	70	90	3.3	-16.7
MW05	5/23/2016	2	90	90	72.1	71.9	70	90	2.1	-18.0
MW06	10/6/2016	2	100	100	86.5	86.3	80	100	6.3	-13.7
MW07	10/10/2016	2	110	110	90.7	90.3	100	110	-9.7	-19.7

Notes:

in = inches; ft = feet; ft amsl = feet above mean sea level; ft bgs = feet below ground surface

Table 2Groundwater Level MeasurementsCity of Issaquah Hydrogeologic Assessment

Well					TOC	Depth to	Groundwater
Owner /	Well	Measurement		Measuring	Elevation	Water	Elevation
Property	Name	Date	Event	Point	(ft amsl)	(ft)	(ft amsl)
		5/17/2016	Baseline	TOC1	<u>81 0</u>	12 /	68 5
COI	MW-1	3/1//2010	sampling	1001	01.9	13.4	08.5
		5/17/2016	Baseline	TOCI	81.8	14 3	67.6
COI	TW-3	5/1//2010	sampling	1001	01.0	11.5	07.0
		5/10/2016	Baseline	TOC1	59.4	8.2	51.2
SPWSD	VT-2.1		sampling		• • • •		
anuan		5/10/2016	Baseline	TOC1	61.8	3.8	58.0
SPWSD	VT-2.2		sampling				
CDWCD	VT 22	5/10/2016	Baseline	TOC1	62.0	3.0	59.0
SPWSD	V 1-2.3		Sampling				
SPWSD	Woll 7_3	5/10/2016	Baseline	Metal rim1	72.1	9.5	62.6
51 (15)	Wen 7-5		Post-				
COI	MW01	6/7/2016	installation	TOC	58.4	3.5	54.9
0.01			Post-				
COI	MW02	6/7/2016	installation	TOC	62.8	1.8	61.0
		(17/2016	Post-	тос	(2.0	1.0	(1.0
COI	MW03	6/ //2016	installation	IUC	62.9	1.0	61.9
		6/7/2016	Post-	TOC	73 1	10.0	63 1
COI	MW04	0///2010	installation	100	75.1	10.0	00.1
COL		6/7/2016	Post-	TOC	71.9	7.8	64.1
	MW05	10/17/2016	installation	TOC	50.4	2.2	55.2
		10/1//2016	Sampling		58.4 (2.8)	3.2	55.2
		10/1//2016	Sampling		02.8	<u> </u>	59.0
		10/1//2016	Sampling	TOC	62.9 72.1	4.1	58.9
		10/1//2016	Sampling	TOC	/ 5.1	/.8	65.2
		10/17/2016	Sampling	TOC	/1.9		03.2 66.0
		10/17/2016	Sampling		<u>80.3</u>	1/.4	72.0
COI		10/1//2016	Sampling	IUC	90.5	18.4	/2.0

Notes:

TOC - Top of casing

ft amsl - feet above mean sea level

¹ Calculated values, well was not surveyed.

Table 3 Summary Water Quality Analytical Results City of Issaquah Hydrogeologic Assessment

Well Owner/	Well	Sampling	6 . I.N	Alkalinty	Calcium	chloride	Magnesium	pH	Potassium	Sodium	Specific Conductance	Sulfate	Total Nitrate	Benzene	Ethyl Benzene	m,p-Xylenes	o-Xylene	Total Xylenes	Methyl-Tert-butyl ethe	Toulene
Property	Name	Date	Sample Name	-	Ca	CI	Mg	pН	K	Na	-	SO ₄	NO ₃				VOCs			
			Units	mg/L as CaCO3	mg/L	mg/L	mg/L	SU	mg/L	mg/L	µmhos/ cm	mg/L	mg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L
COI	COI-5	7/6/2016	COI-Well 5-20160716		34.0	5.30	4.60		3.70	28.0		43.0	< 0.440							
COI	COI-5	7/13/2016	COI-Well5-20160713		36.0	5.10	4.60		4.00	29.0		40.0	$< 0.440 \; J$							
COI	COI-50BS	7/6/2016	COI-Well 50BS-20160716		18.0	3.50	3.50		3.50	19.0		< 0.500	< 0.440							
COI	MW01	6/7/2016	COI-MW01-060716	100	16.0	2.70	6.80	7.9	2.20	22.0	220	3.00	1.30	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5
COI	MW01	10/17/2016	COI-MW01-20161017		20.0	2.80	8.20		2.50	14.0		1.90	$< 0.440 \; J$							
COI	MW02	6/7/2016	COI-MW02-060716	75.0	13.0	2.30	7.30	8.1	1.60	10.0	170	6.40	< 0.440	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5
COI	MW02	10/17/2016	COI-MW02-2016017		20.0	3.50	11.0		1.50	8.80		8.80	1.20							
COI	MW03	6/7/2016	COI-MW03-060716	130	28.0	3.40	13.0	7.7	2.00	11.0	290	11.0	< 0.440	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5
COI	MW03	7/6/2016	COI-MW03-20160716		19.0	3.10	9.00		1.60	12.0		2.00	< 0.440							
COI	MW03	7/13/2016	COI-MW03-20160713		20.0	3.10	10.0		1.40	11.0		2.40	< 0.440							
COI	MW03	7/20/2016	COI-MW03-20160720		21.0	3.20	10.0		1.50	10.0		2.60	< 0.440							
COI	MW03	10/17/2016	COI-MW03-2016017		20.0	3.20	9.60		1.80	9.50		2.10	< 0.440							
COI	MW04	6/7/2016	COI-MW04-060716	61.0	14.0	3.00	5.10	7.2	1.20	7.30	150	4.70	2.60	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5
COI	MW04	7/6/2016	COI-MW04-20160716		18.0	3.70	7.70		1.40	9.40		2.30	1.40							
COI	MW04	7/13/2016	COI-MW04-20160713		22.0	4.00	9.70		1.60	12.0		< 0.500	< 0.440							
COI	MW04	7/20/2016	COI-MW04-20160720		18.0	3.80	7.80		1.20	9.60		2.30	1.40							
COI	MW04	10/17/2016	COI-MW04-20161017		16.0	3.50	6.60		1.60	8.80		4.10	1.60							
COI	MW05	6/7/2016	COI-MW05-060716	58.0	15.0	3.30	5.90	6.6	1.20	7.40	160	8.40	4.60	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5	< 0.5
COI	MW05	7/6/2016	COI-MW05-20160716		14.0	4.30	5.80		< 1.00	7.30		8.20	5.30							
COI	MW05	7/13/2016	COI-MW05-20160713		16.0	3.40	6.20		1.10	8.40		8.30	5.00							
COI	MW05	7/20/2016	COI-MW05-20160720		16.0	3.60	6.50		< 1.00	8.30		8.60	5.10							
COI	MW05	10/17/2016	COI-MW05-20161017		15.0	3.50	5.90		1.00	7.80		8.20	5.20							
COI	MW06	10/17/2016	COI-MW06-20161017		19.0	4.30	8.10		1.80	12.0		8.70	1.10							
COI	MW07	10/17/2016	COI-MW07-20161017		25.0	5.60	13.0		4.90	24.0		11.0	< 0.440							1

Notes:

mg/Lmilligrams per literμg/Lmicrograms per literCaCO3calcium carbonateSUstandard unitsμmhos/cmmicromhos per centimeter

Table 4 Summary of PFC Analytical Results Groundwater and Surface Water City of Issaquah Hydrogeologic Assessment

										р				
			cid				Icid			aci		pi		
			ica		icid	p	nic a	9	р	nic	id	ac	cid	icid
			fon	ació	ic a	aci	fon	acio	aci	lfo	ac	ioi	c a	ic a
			sul	oic	ano	loic	esul	oic	oic	iesu	loic	scar	anoi	ano
			tane	can	dec	otar	Kan	Kan	nan	tar	tar	rade	leci	dec
			phut	oqee	op	hel	her	ohex	IOUG	000	000	oteti	otric	uno
			torc	lorc	lorc	lorc	lorc	lorc	lorc	nor	nor	lorc	lorc	lorc
			rflu	arflu	arth.	ırflu	urflu	arflu	arflı	ll 1	arf)	arflı	irflu	arflu
W-ll Oran and Dara and	Well Nerver	Samula Data Samula Nama		e E		e e	e e e	e e e		ă NEOS	<u>ď</u>	e e e		
well Owner/Property	well Name	Sampling Date Sample Name	PFBS	PFDA	PFDoA	РЕНрА	PFHXS	РГНХА	PFNA	PFOS	PFOA	PFIA	PFIrDA	PFUNA
City of Issamuch	COI6 TMW1	Uni 5/17/2016 COL MW1 051716	$\mu g/L$	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L
City of Issaquah	COLTW3	5/17/2016 COL-TW3-051716	< 0.00300	< 0.00300	< 0.00300	< 0.00300	< 0.00300	< 0.00300	< 0.00300	0.003	< 0.00300	< 0.00300	< 0.00300	< 0.00300
Darigold	ABV249	5/4/2016 Darigold_ABV249-050416	0.00300	< 0.00300	< 0.00300	< 0.00300	< 0.00300	< 0.00300	< 0.00300	0.00300	< 0.00300	< 0.00300	< 0.00300	< 0.00300
Darigold	ABV249	5/4/2016 Darigold-ABV249-050416-DUP	0.0032	< 0.00250	< 0.00250	< 0.00250	0.009	< 0.00250	< 0.00250	0.000	< 0.00250	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW01	6/7/2016 COL-MW01-060716	0.0032	< 0.00250	< 0.00250	< 0.00250	< 0.003	< 0.00250	< 0.00250	< 0.007	0.00230	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW01	10/17/2016 COL-MW01-20161017	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.0008	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW01	6/7/2016 COL-MW01-20101017	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW02	10/17/2016 COL-MW02-2016017	0.00230	< 0.00250	< 0.00250	< 0.00250	0.00230	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW02	6/7/2016 COL-MW03-060716	0.0030	< 0.00250	< 0.00250	0.0230	0.005	0.065	0.0510	0.00230	0.0230	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	6/7/2016 COL-MW03-060716-DUP	0.0920	< 0.00250	< 0.00250	0.029	0.280	0.005	0.0510	0.440	0.0290	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	6/21/2016 COL-MW03-20160621	0.0940	< 0.00250	< 0.00250	0.029	0.300	0.007	0.0310	0.400	0.0300	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	6/21/2016 COL-MW03-20160621-DUP	0.0590	< 0.00250	< 0.00250	0.014	0.170	0.033	0.0180	0.200	0.0200	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	6/28/2016 COLMW03-20160628	0.0490	< 0.00250	< 0.00250	0.013	0.110	0.031	0.0160	0.170	0.0120	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	6/28/2016 COLMW03-20160628-DUP	0.0470	< 0.00250	< 0.00250	0.014	0.097	0.031	0.0130	0.170	0.0120	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	7/6/2016 COL-MW03-20160716	0.0250	< 0.00250	< 0.00250	0.0011	0.057	0.020	0.0061	0.100	0.00510 I	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	7/6/2016 COL-MW03-20160716-Dup	0.0250	< 0.00250 I	< 0.00250	0.0062	0.062	0.015	0.0056	0.100	0.00710 J	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	7/13/2016 COLMW03-20160713	0.0200	< 0.00250 j	< 0.00250	0.0002	0.002	0.014	0.0050	0.100	0.00710 3	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	7/13/2016 COLMW03-20160713 DUP	0.0290	< 0.00250	< 0.00250	0.0071	0.075	0.010	0.0005	0.088	0.0053	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	7/20/2016 COL-MW03-20160720	0.0320	< 0.00250	< 0.00250	0.0004	0.070	0.014	0.0055	0.100	0.0055	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	7/20/2016 COLMW03-20160720-DUP	0.0320	< 0.00250	< 0.00250	0.0071	0.030	0.017	0.0003	0.078	0.0063	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	7/28/2016 COLMW03-20160728	0.0330	< 0.00250	< 0.00250	0.0073	0.170	0.017	0.0250	0.100	0.0005	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	7/28/2016 COLMW03-20160728-DUP	0.0720	< 0.00250	< 0.00250	0.017	0.170	0.035	0.0230	0.330	0.0150	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	10/17/2016 COLMW03-2016017	0.0320	< 0.00250	< 0.00250	0.015	0.160	0.035	0.0025	0.088	0.0150	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW03	10/17/2016 COL-MW03-20161017-DUP	0.0320	< 0.00250	< 0.00250	0.0069	0.007	0.018	0.0055	0.000	0.005	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW04	6/7/2016 COL-MW04-060716	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	0.003	< 0.00250	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW04	6/21/2016 COL-MW04-20160621	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW04	6/28/2016 COI-MW04-20160628	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250
City of Issaquah	MW04	7/6/2016 COI-MW04-20160716	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW04	7/13/2016 COI-MW04-20160713	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW04	7/20/2016 COI-MW04-20160720	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW04	7/27/2016 COI-MW04-20160727	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	0.0039	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW04	10/17/2016 COI-MW04-20161017	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	0.003	< 0.00250	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW05	6/7/2016 COI-MW05-060716	0.0570	< 0.00250	< 0.00250	0.018	0.160	0.040	0.0086	0.390	0.017	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW05	6/21/2016 COI-MW05-20160621	0.0540	< 0.00250	< 0.00250	0.019	0.170	0.042	0.0080	0.490	0.013	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW05	6/28/2016 COI-MW05-20160628	0.0700	< 0.00250	< 0.00250	0.027	0,180	0.056	0.0100	0.500	0.017	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW05	7/6/2016 COI-MW05-20160716	0.056	< 0.00250	< 0.00250	0.017	0.1700	0.039	0.0075	0.4800	0.011	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW05	7/13/2016 COI-MW05-20160713	0.0580	< 0.00250	< 0.00250	0.0250	0.210	0.0540	0.0120	0.440	0.018	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW05	7/20/2016 COI-MW05-20160720	0.052	< 0.00250	< 0.00250	0.019	0.180	0.040	0.0093	0.510	0.013	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW05	7/28/2016 COI-MW05-20160728	0.066	< 0.00250	< 0.00250	0.023	0.18	0.049	0.0087	0.5100	0.018	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW05	10/17/2016 COI-MW05-20161017	0.061	< 0.00250	< 0.00250	0.021	0.17	0.048	0.0079	0.4	0.015	< 0.00250	< 0.00250	< 0.00250
City of Issaguah	MW06	10/17/2016 COI-MW06-20161017	0.096	0.0029	< 0.00250	0.073	0.49	0.22	0.053	2.2	0.08	< 0.00250	< 0.00250	0.016
City of Issaquah	MW07	10/17/2016 COI-MW07-20161017	< 0.00250	< 0.00250	< 0.00250	< 0.00250	0.0049	< 0.00250	< 0.00250	0.0049	0.003	< 0.00250	< 0.00250	< 0.00250

Table 4 Summary of PFC Analytical Results Groundwater and Surface Water City of Issaquah Hydrogeologic Assessment

				Perfluorobutanesulfonic acid	Perfluorodecanoic acid	Perfluorododecanoic acid	Perfluoroheptanoic acid	Perfluorohexanesulfonic acid	Perfluorohexanoic acid	Perfluorononanoic acid	Perfluorooctanesulfonic acid	Perfluorooctanoic acid	Perfluorotetradecanoic acid	Perfluorotridecanoic acid	Perfluoroundecanoic acid
Well Owner/Property	Well Name	Sampling Date	Sample Name	PFBS	PFDA	PFDoA	PFHpA	PFHxS	PFHxA	PFNA	PFOS	PFOA	PFTA	PFTrDA	PFUnA
Sammamish Plateau Water Sewer District	SPWSD-VT2.1	5/10/2016	SPWSD-VT2.1-051016	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250
Sammamish Plateau Water Sewer District	SPWSD-VT2.2	5/10/2016	SPWSD-VT2.2-051016	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250
Sammamish Plateau Water Sewer District	SPWSD-VT2.3	5/10/2016	SPWSD-VT2.3-051016	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250
Sammamish Plateau Water Sewer District	SPWSD-Well7	6/15/2016	S-06	< 0.0900	NA	NA	< 0.0100	0.0140 J	NA	< 0.0200	0.0190 J	< 0.0200	NA	NA	NA
Sammamish Plateau Water Sewer District	SPWSD-Well7	7/26/2016	Well 7 (201607270134)	0.0043	< 0.00250	< 0.00250	< 0.00250	0.012	0.0026	0.0028	0.015	< 0.00250	< 0.00250	< 0.00250	< 0.00250
Sammamish Plateau Water Sewer District	SPWSD-Well7.3	5/10/2016	SPWSD-Well7.3-051016	0.0054	< 0.00250	< 0.00250	< 0.00250	0.019	< 0.00250	0.0034	0.019	< 0.00250	< 0.00250	< 0.00250	< 0.00250
Sammamish Plateau Water Sewer District	SPWSD-Well8	7/26/2016	Well 8 - Sample 1 (201607270135)	0.0047	< 0.00250	< 0.00250	< 0.00250	0.019	< 0.00250	0.0056	0.026	< 0.00250	< 0.00250	< 0.00250	< 0.00250
Sammamish Plateau Water Sewer District	SPWSD-Well8	7/26/2016	Well 8 - Sample 2 (201607270137)	0.005	< 0.00250	< 0.00250	< 0.00250	0.020	< 0.00250	0.006	0.029	< 0.00250	< 0.00250	< 0.00250	< 0.00250
Sammamish Plateau Water Sewer District	SPWSD-Well9	6/20/2016	S-13	< 0.0900	NA	NA	< 0.0100	< 0.0300	NA	< 0.0200	< 0.0400	< 0.0200	NA	NA	NA
Surface water samples, Issaquah Creek	COI-SW-A	9/27/2016	COI-SW-A-092716	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250
Surface water samples, Issaquah Creek	COI-SW-B	9/27/2016	COI-SW-B-092716	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250

Notes:

Shaded - Value detected above laboratory reporting limit.

Bold - Bold text indicates the results for groundwater are above the corresponding US EPA Provisional Health Advisory Levels of 0.07 ug/L for PFOS and 0.07 ug/L for PFOA or 0.07 for combined PFOS and PFOA concentrations in groundwater (May 2016).

COI City of Issaquah

* City monitoring well COI-MW1 is renamed to COI6-TMW1s to differentiate from new monitoring well MW01.

** Sample analyzed by EPA Method 537 at Anatek Labs Inc, Moscow, ID.

- µg/L micrograms per liter
 - < Value is less than the laboratory reporting limit shown
- J Estimated value
- NA Not analyzed

Samples analyzed by EPA Method 537 at Eurofins Eaton Analytical Laboratory in Monrovia, CA, unless otherwise noted.

COI-MW-20160716 sample names: the date was incorrectly noted in the field name, the samples were collected on 07/06/16 as indicated by the sample date.

Analytical results were reviewed and validated (Stage 2A) in accordance with guidelines provided in the EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review, August 2014 (USEPA-540-R-014-002). Full validation memoranda and analytical laboratory reports are provided as attachments to this technical memorandum.

Sammamish Plateau Sewer and Water District (SPSWD) Wells 7, 8, 9 were sampled by SPSWD on the dates shown; these results are included for reference only. These results were not validated by the COI.

Table 5 Summary of PFCs Analytical Results at 175 Newport Way NW City of Issaquah Hydrogeologic Assessment

				Perfluorobutanesulfonic acid	Perfluorodecanoic acid	Perfluorododecanoic acid	Perfluoroheptanoic acid	Perfluorohexanesulfonic acid	Perfluorohexanoic acid	Perfluorononanoic acid	Perfluorooctanesulfonic acic	Perfluorooctanoic acid	Perfluorotetradecanoic acid	Perfluorotridecanoic acid	Perfluoroundecanoic acid
_			Soil Sample Depth												
Property	Sample Name	Date Sampled	(inches bgs)	PFBS	PFDA	PFDoA	PFHpA	PFHxS	PFHxA	PFNA	PFOS	PFOA	PFTA	PFTrDA	PFUnA
			Units	ng/g	ng/g	ng/g	ng/g	ng/g	ng/g	ng/g	ng/g	ng/g	ng/g	ng/g	ng/g
175 Newport Way NW	COI-STSP01-20160722	2016-07-22	26.5-27	< 1.90	1.2	< 0.960	< 0.720	< 1.90	0.58	0.77	23	< 0.720	< 0.960	< 1.40	1.0
175 Newport Way NW	COI-STTA01-20160722	2016-07-22	26.5-27	8.9	< 0.540	< 1.10	5.1	80	30	9.7	1,300	11	< 1.10	< 1.60	2.8
175 Newport Way NW	COI-STTA02-20160722	2016-07-22	46-46.5	4.5	3.9	< 1.10	2.1	25	15	33	180	4.3	< 1.10	2.7	36 J
175 Newport Way NW	COI-STTA02-20160722-Dup	2016-07-22	46-46.5	5.4	4.5	< 1.20	2.4	29	19	43	250	5.2	< 1.20	3	63 J

Notes:

Shaded - Value detected above laboratory reporting limit.

mg/kg milligrams per kilogram

ng/g nanogram per gram

< Value is less than the laboratory reporting limit shown

J Estimated value

bgs below ground surface

Samples analyzed by EPA Method 537 at Eurofins Eaton Analytical Laboratory in Monrovia, CA, unless otherwise noted.

Analytical results were reviewed and validated (Stage 2A) in accordance with guidelines provided in the EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review, August 2014 (USEPA-540-R-014-002). Full validation memoranda and analytical laboratory reports are provided as attachments to this technical memorandum.

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FIGURES





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Notes:	
Qa - Quaternary Alluvium	
Qvi - Quaternary Ice Contact Deposits	
Qva - Quaternary Advance Outwash Deposits	
20x Vertical Exaggeration	
*Wells Projected into line of section	
(XX Ft) - Ground Elevation of Well	





Leger	nd Existing Well Newly Installed Monitoring Well Groundwater Elevation	Notes 1. Aerial image from 2012. 2. Groundwater Elevations in feet above mean sea level. N	Groundwater Iss	Elevations, October aquah, Washington	2016
->	Estimated Groundwater Flow Direction	Ĩ	Geosy	mtec <a>>	Figure
(54.88)	- Groundwater Elevation (hand measured in October 2016).	0 0.5 Miles	Seattle, Washington	November 2016	3

\\SantaBarbara-01\Data\GIS\PNG0714 - Issaquah\Projects\COI_Report\Fig03_GW_Levels.mxd 11/18/2016 3:22:15 PM















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APPENDIX A DISCRETE DEPTH GROUNDWATER SAMPLING DURING DRILLING

1. DISCRETE DEPTH GROUNDWATER SAMPLING DURING DRILLING

Discrete-depth groundwater samples were obtained during drilling as described in the Work Plan (Geosyntec, 2016). The methodology for sampling during drilling was as follows:

- MW01 samples were obtained using disposable bailers for the shallow and deep samples. A second discrete-depth groundwater sample was obtained at MW01 at the total borehole depth of 80 feet bgs. Approximately one casing volume of water was purged prior to collecting the deeper groundwater sample because water was added during drilling (due to heaving sand conditions).
- MW02, MW04, and MW05 a peristaltic pump and new, disposable low-density polyethylene (LDPE) tubing were used to purge the borehole and sample first encountered groundwater.
- MW03 was purged and sampled at first encountered groundwater using a peristaltic pump and new, disposable LDPE tubing. A second sample was collected at approximately 40 to 50 feet bgs, and three casing volumes of water were purged prior to sampling because water was added during drilling (due to heaving sand conditions). A stainless steel Monsoon® pump and disposable LDPE tubing were used for the purging and sampling at the deeper depth. An equipment blank was collected from the Monsoon® pump and tubing before purging.
- MW06 and MW07 a peristaltic pump and new, disposable LDPE tubing were used to purge the borehole and sample first encountered groundwater. Groundwater samples were also obtained at MW06 at depths of 34.5 to 39.5 and 51 to 56, feet bgs, and at MW07 at 35 to 40 and 65 to 70 feet bgs.

Groundwater samples were placed directly into laboratory supplied-containers, placed into a cooler with ice, and submitted for analytical testing on a standard turn-around time. Each cooler was shipped to the laboratory with a trip blank. Thirteen groundwater samples, five trip blanks, and one equipment blank were analyzed by Eurofins Eaton Analytical Laboratory in Monrovia, CA for PFCs using Environmental Protection Agency (EPA) Method 537.

1.1 Discrete-Depth Analytical Results

PFCs were detected in nine groundwater samples from MW02, MW03, MW04, MW05, MW06, and MW07 as follows:

• MW01: PFCs were not detected (<0.0025 micrograms per liter $[\mu g/L]$) in either the shallow (10 to 20 feet bgs) or deep (55 to 65 feet bgs) samples.

- MW02: PFOS and PFOA were not detected (<0.0025 μg/L) in the sample from 40 to 50 feet bgs, although perflourobutanesulfonic acid (PFBS), perfluorohexanesulfonic acid (PFHxS), and perfluorohexanoic acid (PFHxA) were detected.
- MW03: PFOS was detected at 0.01 and 0.11 μ g/L and PFOA at 0.0046 and 0.0075 μ g/L in the samples from 10 to 20 and 40 to 50 feet bgs, respectively. Other PFCs detected in both groundwater samples included PFBS, PBHpA, PFHxS, PFHxA, and PFNA. The PFOS concentration in the deeper sample was an order of magnitude higher than the shallow sample.
- MW04: PFOS was detected at a concentration of 0.003 μ g/L, and PFOA was not detected (<0.0025 μ g/L). PFBS and PFHxS were also detected in the sample from MW04.
- MW05: PFOS and PFOA were detected at 0.013 µg/L at 0.022 µg/L, respectively. Other PFCs detected in MW05 included PFBS, PBHpA, PFHxS, PFHxA, and PFNA.
- MW06: PFOS was detected at 0.30 μg/L, 0.50 μg/L, 0.74 μg/L and PFOA was detected 0.021 μg/L, 0.036 μg/L, 0.040 μg/L (with increasing depth, shallow to deepest). Other PFCs detected in MW06 included PFBS, PBHpA, PFHxS, PFHxA, PFNA, and Perfluoroundecanoic acid (PFUnA).
- MW07: PFOS and PFOA were detected in the shallowest sample at 0.025 and 0.029 μ g/L. PFOS was detected at 0.004 and 0.018 μ g/L at the next two depths sampled. Other PFCs detected in MW07 included PFBS, PBHpA, PFHxS, and PFHxA.

The EPA provisional health advisory level for PFOS and PFOA, individually or summed together, is 0.07 μ g/L. The PFOS concentration (0.11 μ g/L) in one sample from MW03 (40 to 50 feet bgs) and the three samples at MW06 (ranging from 0.3 to 0.74 μ g/L) were greater than the EPA provisional health advisory level. PFOS concentrations were below the EPA level at MW02, MW03 (10 to 20 feet bgs), MW04, MW05, and MW07. PFOA was detected below the EPA level at MW03 (2 samples), MW05, and MW07 (1 sample).

Analytical results are summarized in Table A-1. Analytical laboratory reports are included in Attachment B along with a Data Quality Review Memo. Results of the data quality review indicate that analytical data are acceptable.

Table A-1 Summary of Discrete Depth Sample PFCs Analytical Results City of Issaquah Hydrogeologic Assessment

Well Owner/		Sampling		Perfluorobutanesulfonic acid	Perfluorodecanoic acid	Perfluorododecanoic acid	Perfluoroheptanoic acid	Perfluorohexanesulfonic acid	Perfluorohexanoic acid	Perfluorononanoic acid	Perfluorooctanesulfonic acid	Perfluorooctanoic acid	Perfluorotetradecanoic acid	Perfluorotridecanoic acid	Perfluoroundecanoic acid
Property	Well Name	Date	Sample Name	PFBS	PFDA	PFDoA	PFHpA	PFHxS	PFHxA	PFNA	PFOS	PFOA	PFTA	PFTrDA	PFUnA
			Units	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L	μg/L
					Discrete D	epth Samp	ling Result	S							
COI	MW01	5/26/2016	MW01_30to40_20160526	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025
COI	MW01	5/26/2016	MW01_55to65_20160526	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025
COI	MW02	5/31/2016	MW02_40 to 50_20160531	0.0073	< 0.0025	< 0.0025	< 0.0025	0.0077	0.0051	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025	< 0.0025
COI	MW03	5/24/2016	MW03_10 to 20_20160524	0.019	< 0.0025	< 0.0025	0.0051	0.031	0.012	0.0027	0.01	0.0046	< 0.0025	< 0.0025	< 0.0025
COI	MW03	5/23/2016	MW03_40 to 50_20160524	0.036	< 0.0025	< 0.0025	0.0086	0.08	0.021	0.0091	0.11	0.0075	< 0.0025	< 0.0025	< 0.0025
COI	MW04	5/27/2016	MW04_29to39_20160527	0.0051	< 0.0025	< 0.0025	< 0.0025	0.0084 U	< 0.0025	< 0.0025	0.0028 U	< 0.0025	< 0.0025	< 0.0025	< 0.0025
COI	MW05	5/23/2016	MW05_10 to 20_20160523	0.03	< 0.0025	< 0.0025	0.016	0.039	0.036	0.0084 U	0.013	0.022	< 0.0025	< 0.0025	< 0.0025
COI	MW06	10/5/2016	COI-MW06-20161005-19.5-24.	0.03	< 0.00250	< 0.00250	0.026	0.14	0.061	0.0069	0.3	0.021	< 0.00250	< 0.00250	< 0.00250
COI	MW06	10/6/2016	COI-MW06-20161006-34.5-39.	0.052	< 0.00250	< 0.00250	0.044	0.24	0.1	0.021	0.5	0.036	< 0.00250	< 0.00250	0.0062
COI	MW06	10/6/2016	COI-MW06-20161006-51-56	0.073	< 0.00250	< 0.00250	0.05	0.3	0.13	0.022	0.74	0.040	< 0.00250	< 0.00250	0.019
COI	MW07	10/7/2016	COI-MW07-20161007-20-25	0.0044	< 0.00250	< 0.00250	0.0076	0.0094	0.011	< 0.00250	0.03	0.029	< 0.00250	< 0.00250	< 0.00250
COI	MW07	10/7/2016	COI-MW07-20161007-35-40	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	< 0.00250	0.0	< 0.00250	< 0.00250	< 0.00250	< 0.00250
COI	MW07	10/7/2016	COI-MW07-20161007-65-70	< 0.00250	< 0.00250	< 0.00250	< 0.00250	0.004	< 0.00250	< 0.00250	0.0	< 0.00250	< 0.00250	< 0.00250	< 0.00250

Notes:

Bold

Shaded - Value detected above laboratory reporting limit.

Values are above the corresponding US EPA Provisional Health Advisory Levels of 0.07 ug/L for PFOS and 0.07 ug/L for PFOA or 0.07 for combined PFOS and PFOA concentrations.

APPENDIX B BOREHOLE AND MONITORING WELL CONSTRUCTION LOGS



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PNG0714 PORTLAND



GLB **IBRARY** GINT DEFAULT EED ISSAQUAH.GPJ

PNG0714 PORTLAND

Ge	consult	621 SW Morris Portland, Oreg Phone: 503.22	son Street on 97205 22.9518	, Suite	e 600	BORING START FINISH PROJEG LOCAT	G DATE DATE CT P ION Is	MW01 5/26/2016 5/26/2016 hase 2 Hydrogeo saquah, Washing	logic gton	Elevat Invest	tion igatio	SHE 58.36 F1 n	et 1 of 3 T. MSL
	DRE3 10/00			JG		PROJE		MBER PNG0714				1]
DEPTH (ft)		MATERIAL DESCRIPTION	SYMBOLIC LOG	MELL LOG	W CONST MAT	ell Ruction Erial	ELEVATION (ft)	SAMI SAMPLE NAME	JES	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	Grass, topsoil Brown (7.5YR 4, graded, some ro	/3), loose, dry, silty SAND; well unded gravel			Surface complet Morris monume 2" sch 4 well cas	ion: 8" ent 0 PVC ing	- 55 - - 50 - - - - - - - - - - - - - -			40	0.1	SM	
15 — - - - 20 — - - - - -	Brown (7.5YR 4, dry, sandy SILT; gravel Gray (Gley 2 4/5 sandy SILT	/3), medium dense, moist to ; some gray mottling, trace fine 5B dark bluish gray), firm, wet,			Hallibur Hole-plu bentonit	on Ig, 3/8" e chips	- 40 - - - - 35 -			35		SM-ML	
25 — — 30 — — 25	6 fine sand trace organic Gray (Gley 2 4/5 fine to medium s graded, trace roo	s/wood fragments iB dark bluish gray), loose, wet SAND; well sorted, poorly unded gravel			2" sch 4 0.01" sla screen	0 PVC otted well	30 - - - - - - - - - - - - - - - - - - -	MW01_30to40 _20160526 _@1215		100	0.1	SM	no oxidation noted First encountered groundwater: 29' DTW: 18' Screen: 30-40'
CONT EQUIF DRILL DIAMI	RACTOR Holt Services NORTHING 201387.690 PMENT Terra Sonic, track EASTING 1338963.340 MTHD Sonic ANGLE ETER 8" BEARING BEARING GER J Dahl REVIEWER C Bartlett											5	

Ge	Consultants Consultants Consultants CORM: CORE: 10/00 CONSULTANCE	rrison Street, egon 97205 222.9518 IOLE LC	Suite 60	00	BORING START FINISH PROJEC LOCAT PROJEC	GATE DATE DATE CT P ION Is CT NUM	MW01 5/26/2016 5/26/2016 hase 2 Hydrogeo saquah, Washing IBER PNG0714	logic gton	Elevat Invest	ion igatio	SHE 58.36 F1 n	ET 2 OF 3 r. MSL
DEPTH (ft)	MATERIAL DESCRIPTION	SYMBOLIC LOG	MELL LOG	WI CONSTR MATI	ELL RUCTION ERIAL	ELEVATION (ft)	SAMI SAMPLE NAME	TYPE	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	Gray (Gley 2 4/5B dark bluish gray), firm, we sandy SILT trace organics/wood fragments firm to stiff organic odor Gray (Gley 2 4/5B dark bluish gray), firm, we sandy SILT; trace organics and wood fragments Gray (Gley 2 4/5B dark bluish gray), medium dense, wet, fine to medium silty SAND Gray (Gley 2 4/5B dark bluish gray), firm, we sandy SILT; trace organics and wood fragments	t,		Halliburt Hole-plu bentonite	on g, 3/8" e chips		MW01_55to65 _20160526 _@1600		90	0.2	SM-ML SM-ML SM-ML	10' of sand heave no oxidation noted DTW: 13' Screen: 55-65' 15' of sand heave
70 - Cont Equif Drill Dami Logg	Gray (Gley 2 4/5B dark bluish gray), medium dense, wet, fine to medium silty SAND; well sorted DNTRACTOR Holt Services NORTHING 201387.690 RUIPMENT Terra Sonic, track RILL MTHD Sonic ANGLE Vertical BEARING BEARING SOUCH ANGLE Vertical BEARING BEARING SOUCH ANGLE Vertical BEARING SOUCH ANGLE Vertical BEARING SOUCH ANGLE Vertical BEARING SOUCH ANGLE VERTICAL BEARING SOUCH ANGLE VERTICAL SOUCH ANGLE VERTICAL SOUCH ANGLE VERTICAL SOUCH ANGLE VERTICAL SOUCH ANGLE VERTICAL											

Ge	COSY cons		eco nts BORE	orrison Dregon 3.222.9	Street, 97205 9518	Suite 6	00	BORING START FINISH PROJEC LOCATI PROJEC	GATE DATE DATE CT P ION IS	MW01 5/26/2016 5/26/2016 Phase 2 Hydrogeo ssaquah, Washing MBER PNG0714	logic gton	Elevat Invest	ion igatio	SHE 58.36 F1 n	iet 3 of 3 T. MSL
DEPTH (ft)		D	MATERIAL ESCRIPTION		SYMBOLIC LOG	MELL LOG	WE CONSTF MATE	ELL RUCTION ERIAL	ELEVATION (ft)	SAMIPLE NAME	PLES	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	Gray (Gley dense, wet well sorted Gray (Gley moist, SILT Gray (Gley dense, wet sorted	2 4/5B , fine to 2 4/5B T; non-p 2 4/5B t, fine to	dark bluish gray), mediun medium SAND; trace gr dark bluish gray), firm, plastic dark bluish gray), mediun medium silty SAND; wel	m ; avel, ; m • II					- -15 - - - -20 - - 20 - - - 20 -			90	0.4	SW ML SM	
CONT EQUIP DRILL DIAME LOGG	RACTOR PMENT MTHD ETER GER J Dahl	Holt S Terra Sonic 8"	Services Sonic, track REVIEWER C Bartlett	NOR EAS ANG BEA PF	THING TING GLE RING RINTED	20138 13389 Vertic 07/08	87.690 963.340 eal 8/16	COORD	KS: Si INATE	te: park bench. We SYSTEM: DR SYMBOLS AND ABB	ell Tag REVIATI) ID: B. IONS	IX-185		

Ge	Consultants BOREN BOREL 621 SW Mo Portland, Ol Phone: 503 BOREL	rrison Str egon 972 222.9518 IOLE	reet, 205 8 LO	Suite 6	00	BORING START FINISH PROJEG LOCAT	G DATE DATE CT P ION Is CT NUM	MW02 5/31/2016 6/2/2016 hase 2 Hydrogeo ssaquah, Washing /IBER PNG0714	logic gton	Elevat	tion igatio	SHE 59.70 FT n	ET 1 OF 3
DEPTH (ft)	MATERIAL DESCRIPTION		STMBULIC LUG	MELL LOG	WI CONSTF MATI	ELL RUCTION ERIAL	ELEVATION (ft)	SAMI SAMPLE NAME	PLES	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
5	Topsoil Brown (10YR 6/6 brownish yellow), medium stiff, dry, sandy SILT; trace oxidation becomes moist Brown (10YR 6/6 brownish yellow), medium dense, wet, silty fine SAND; well graded, po sorted, some oxidation Gray (Gley 2 4/5B dark bluish gray), firm, we sandy SILT; trace wood fragments	x 1 kg 			Above gr completi feet), ste monume bollards around monume	round on (+3.5 el nt, 3 installed nt	- - 55_ - - 50_			70		SM-ML SM-ML	
	3" lens of gray, loose, wet, medium sand Gray (Gley 2 4/5B dark bluish gray), stiff, dry SILT; slightly mottled Gray (Gley 2 4/5B dark bluish gray), firm, moist, sandy SILT 3" lens of gray, loose, wet, medium sand becomes wet				2" sch 40 well casi) PVC ng	- 45_ - 40_ -			100		ML SM-ML	
	Gray (Gley 2 4/5B dark bluish gray), firm, moist, sandy SILT trace wood fragments						- 35_ - - 30_ - - - - - 25					ML	
35 CONT EQUIF DRILL DIAMI LOGG	RACTOR Holt Services PMENT Terra Sonic, track MTHD Sonic ETER 8" GER J Dahl REVIEWER C Bartlett	NORTH EASTIN ANGLE BEARIN PRINT	ING IG IG IG TED	2013: 13407 Vertic 07/08	54.260 787.500 cal 3/16	REMAR COORD SEE KEY S	KS: Sit	e: Pickering Trail. SYSTEM: R SYMBOLS AND ABBI	Well	<u>™</u> Tag ID Ions	: BJX-	187	

Ge	Consultants Consu	n Street n 97205 .9518 L E LC	, Suite 60	00 BORING START FINISH PROJEC LOCAT PROJEC	GATE DATE DATE CT P ION IS	MW02 5/31/2016 6/2/2016 Phase 2 Hydrogeo ssaquah, Washing MBER PNG0714	logic I gton	Elevat nvest	tion igatio	SHE 59.70 F1 n	et 2 of 3 T. MSL
DEPTH (ft)	MATERIAL DESCRIPTION	SYMBOLIC LOG	MELL LOG	WELL CONSTRUCTION MATERIAL	ELEVATION (ft)	SAM SAMPLE NAME	PLES BdL	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	becomes wet becomes moist becomes wet Gray, medium dense, wet, SAND Gray (Gley 2 4/5B dark bluish gray), firm, wet, sandy SILT becomes moist 6" lens gray, medium dense, sand becomes moist to dry, trace wood fragments becomes wet Gray (Gley 2 4/5B dark bluish gray), medium dense, wet, fine to medium sandy GRAVEL; poorly sorted, well graded, trace organics and mottling Gray (Gley 2 4/5B dark bluish gray), firm to soft, wet, sandy SILT; trace wood fragments 2-6" lens wet, fine to medium sand Gray (Gley 2 4/5B dark bluish gray), lirm to soft, wet, sandy SILT; trace wood fragments		20135	Halliburton Hole-plug, 3/8" bentonite chips	- 20_ - 20_ - - - 15_ - - 10_ - - - - - - - - - - - - - - - - - - -	MW02_40to50 _20160531 @1730		100 100	BJX-	SM SM-ML GM SM-ML	First encountered groundwater: 39' DTW: 2.0' Screen: 40-50' End of drilling 05/31/2016; continue on 06/01/2016
EQUIF DRILL DIAMI LOGG	PMENT Terra Sonic, track EA PMENT Terra Sonic, track EA MTHD Sonic AN ETER 8" BE EER J Dahl REVIEWER C Bartlett P	RIHING STING GLE ARING PRINTED	20135 13407 Vertic 	4.200 REMAR 87.500 al /16 COORD SEE KEY S	NATE	SYSTEM: DR SYMBOLS AND ABB	vveii i <u>Reviatio</u>	ag ID:	. BJX-	18/	

(PORTLAND) PNG0714 ISSAQUAH.GPJ EED DEFAULT GINT LIBRARY.GLB 7/8/16 I OG W/WELL SONC <u>c</u>

Ge	Consultants BOREHC BOREHC CONSULTANTS 621 SW Morriss Portland, Orego Phone: 503.222 BOREHC	on Street, on 97205 2.9518	Suite 6	00	BORING START FINISH PROJEG LOCAT PROJEG	DATE DATE DATE CT P ION Is CT NUM	MW02 5/31/2016 6/2/2016 Phase 2 Hydrogeo ssaquah, Washing MBER PNG0714	logic gton	Elevat Invest	tion igatio	SHE 59.70 F1 n	iet 3 of 3 r. MSL
DEPTH (ft)	MATERIAL DESCRIPTION	SYMBOLIC LOG	MELL LOG	W CONSTR MAT	ELL RUCTION ERIAL	ELEVATION (ft)	SAMF SAMFLE NAME	PLES Bd AL	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	medium to coarse SAND; well graded, poorly sorted, some gravel Brown (10YR 5/1 gray), loose, dry, medium to coarse SAND; well graded, poorly sorted, some gravel 6" lens gray, firm, silt Gray (Gley 2 4/5B dark bluish gray), loose, dry, medium to coarse SAND; well graded, poorly sorted, some gravel Gray (Gley 2 4/5B dark bluish gray), medium dense, wet, medium to coarse SAND; poorly graded, no fines, trace coarse gravels 6" lens firm, wet, silt Gray (Gley 2 4/5B dark bluish gray), firm, moist, SILT Gray (Gley 2 4/5B dark bluish gray), medium dense, wet, fine to coarse SAND; well graded Gray (Gley 2 4/5B dark bluish gray), medium dense, wet, silty fine SAND; poorly graded, well sorted			pre-pack	screen	15_ 15_ 15_ 			100		SW SW ML SW-ML	10-15' of sand heave
CONT EQUIF DRILL DIAMI LOGO	RACTORHolt ServicesNOPMENTTerra Sonic, trackE4MTHDSonicA1ETER8"B1GER J DahlREVIEWER C Bartlett	ORTHING ASTING IGLE EARING PRINTED	20135 13407 Vertic 07/08	54.260 787.500 al 9/16	COORD SEE KEY S	KS: Sit	te: Pickering Trail. SYSTEM: DR SYMBOLS AND ABBI	Well ⁻	Tag ID	: BJX-	187	

Ge	consul	tec ^b tants 621 SW Morr Portland, Ore Phone: 503.2 BOREH	600	BORING START FINISH PROJEC LOCAT PROJEC	GATE DATE DATE CT P ION Is CT NUI	MW03 5/24/2016 5/25/2016 Phase 2 Hydrogeo ssaquah, Washing MBER PNG0714	logic gton	Elevat Invest	tion igatio	SHE 63.16 F1 n	iet 1 of 3 T. MSL		
DEPTH (ft)		MATERIAL DESCRIPTION	SYMBOLIC LOG	MELL LOG	W CONST MAT	ELL RUCTION ERIAL	ELEVATION (ft)	SAMPLE NAME	PLES	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	Brown (2.5YR 5 SILT Brown (7.5YR 4 oxidation, loose trace silt Brown (2.5YR 6 widation, very Brown (2.5YR 6 mottles, mediu some oxidation Brown (2.5YR 6 medium to fine graded, poorly Gray (Gley 2 5/ Isilty SAND; poor organics Gray (Gley 2 5/ wet, SILT Gray (Gley 2 5/ wet, SILT Gray (Gley 2 4/ dense, wet, fine sorted, well gra Gray (Gley 2 4/ medium dense, poorly sorted, v Gray (Gley 2 4/ wet, gravelly cc Gray (Gley 2 4/ stiff, wet, SILT;	5/2 weak red), firm, dry, sandy 4/6 strong brown) with trace rec e, wet, fine to medium SAND; 4/6 strong brown) with trace rec stiff, moist, SILT 5/2 weak red) with some gray n dense, moist, sandy SILT; 5/2 weak red), loose, wet, SAND; some oxidation, well sorted 5BG greenish gray), firm, mois orly graded, well sorted, trace 5BG greenish gray), medium y SAND 5BG greenish gray), medium y SAND 5BG greenish gray), wet, stiff, 10B dark bluish gray), medium a to medium SAND; poorly ded 5BG dark greenish gray), wet, fine to coarse SAND; vell graded 5BG dark greenish gray), very slightly plastic, trace fine sand			Surface completi Morris monume 2" sch 4 well cas	on: 8" ent		MW03_10to20 _20160524 @1545		90	0.2 0.2 0.2 0.2 0.2 0.1 0.2 0.1	SM-ML SM ML SM-ML SM ML SW SP ML	DTW: 8.27' Screen: 10-20' Driller using water to drill, heaving conditions
CONT EQUIF DRILL DIAMI LOGG	RACTOR Hol PMENT Ter MTHD Sor ETER 8'' ER J Dahl	It Services rra Sonic, track nic REVIEWER C Bartlett	NORTHING EASTING ANGLE BEARING PRINTEL	6 2000 134 ² Vert	673.280 1038.950 tical - 08/16	COORD SEE KEY S	INATE	te: Dental lab. We SYSTEM: DR SYMBOLS AND ABB	II Tag	ID: BJ)	X-184		

Ge	Consultants BOREI BOREI CONSULTATION CON	rrison Street, regon 97205 .222.9518 HOLE LC	Suite 600	BORING START FINISH PROJEC LOCAT PROJEC	G DATE DATE CT P ION Is CT NUM	MW03 5/24/2016 5/25/2016 Phase 2 Hydroged ssaquah, Washing MBER PNG0714	logic gton	Elevat Invest	tion igatio	SHE 63.16 F⊺ n	iet 2 of 3 r. MSL	
DEPTH (ft)	MATERIAL DESCRIPTION	SYMBOLIC LOG	MELL LOG	WELL ISTRUCTION /ATERIAL	ELEVATION (ft)	SAM SAMPLE NAME	PLES TYPE	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS	
	Gray (Gley 2 4/5BG dark greenish gray), loo wet, fine to coarse SAND; with gravel interbo Gray (Gley 2 4/5BG dark greenish gray), loo wet, gravelly fine to coarse SAND; fine to coarse gravel	Se,	Halli Hole bent (12-1	burton plug, 3/8" onite chips 50lb bags)	- 25 - - - 20 - - -			100		SM		
	Gray (Gley 2 4/5BG dark greenish gray), stif wet, sandy SILT; trace organics	tar (ertar) 1 (ertar) 1.			- 15 ⁻ - - - 10 ⁻ -			100	0	ML	possible heave	
	Gray (Gley 2 4/5B dark bluish gray), dense, wet, silty fine to coarse SAND; trace organic well graded. poorly sorted				- 5 - - - - - - - - - - - - - - - - - -			100	0	SM		
70 - Cont Equif Drill Logg	well graded, poorly sorted well graded, poorly sorted - - - - 0											

Ge	consultants B	1 SW Morrison Street rtland, Oregon 97205 one: 503.222.9518	, Suite 60	00	BORING START FINISH PROJEC LOCATI PROJEC	DATE DATE DATE CT P ION Is CT NUN	MW03 5/24/2016 5/25/2016 hase 2 Hydrogeo saquah, Washing /IBER PNG0714	logic gton	Elevat Invest	ion igatio	SHE 63.16 F1 n	et 3 of 3 T. MSL
DEPTH (ft)	MATERIAL DESCRIPTION	SYMBOLIC LOG	MELL LOG	WEI CONSTRU MATEI	LL UCTION RIAL	ELEVATION (ft)	SAMI SAMPLE NAME	PLES	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	Gray (Gley 2 4/5B dark bluish gray sandy SILT trace oxidation at contact Brown (10YR 4/3), loose, gravelly S to coarse, rounded gravel, well sor graded (Advanced Outwash) medium to coarse medium to coarse Gray (Gley 2 5/5B bluish gray), fine SAND fine to medium End of boring, install monitoring we	e to coarse		2" sch 40 l 0.01" slotte screen	PVC ed well	-10 - -10 - - - -15 - - - - - - - - - - - - - - - - - - -			100 100 90	0.6	ML SP	10' sand heave flowing well, water level fluctuating at ground
	RACTOR Holt Services PMENT Terra Sonic, track MTHD Sonic ETER 8" GER J Dahl REVIEWER C	NORTHING EASTING ANGLE BEARING Bartlett PRINTED	20067 13410 Vertic 0 07/08	73.280 938.950 cal 9/16	REMAR COORD	KS: Sit	e: Dental lab. Wel SYSTEM: IR SYMBOLS AND ABBI	ll Tag REVIAT	ID: BJ)	K-184		

Ge	Consultants BOREHC CONSULTANTS 621 SW Morris Portland, Oreg Phone: 503.22 BOREHC	MW04 5/27/2016 5/31/2016 hase 2 Hydrogeo ssaquah, Washing MBER PNG0714	logic Iton	Elevat Invest	tion igatio	SHE 73.30 FT n	ET 1 OF 3 . MSL					
DEPTH (ft)	MATERIAL DESCRIPTION	SYMBOLIC LOG	MELL LOG	WI CONSTF MATI	ELL RUCTION ERIAL	ELEVATION (ft)	SAMF BWDLE NAME SAMPLE	TYPE	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	Grass, topsoil Brown, medium stiff, dry, sandy SILT; some rounded gravel			Surface completi Morris monume	on: 8" nt	- 70 ⁻ - - 65 ⁻				0.2	ML	
10 — 10 — - - 15 —	Gray, medium dense, dry, medium to fine SAND; poorly graded, trace gravel Gray with some brown, medium dense, dry, silty SAND; poorly graded, trace gravel Brown, firm, wet, SILT; oxidation					- - 60 - -			60	0	SM SM ML	
	Gray (Gley 2 4/5BG dark greenish gray), medium dense, moist, fine silty SAND; poorly graded, trace gravel, oxidation at 16.5-17.0 Gray (Gley 2 4/5BG dark greenish gray), medium dense, fine silty SAND; poorly graded			2" sch 40 well casi) PVC ng	- 55 ⁻ - -			70		SM SM	
 	Gray (Gley 2 4/5BG dark greenish gray), firm, moist, SILT Gray (Gley 2 4/5BG dark greenish gray), firm, dry, SILT; some oxidation Black, dry, PEAT; some wood pieces, blocky structure					- 50 - - - -					ML ML PT	
	6" gray, firm, wet, silt Gray (Gley 2 4/5BG dark greenish gray), loose.					45 - - - 40 - -	MW04_29to39 _20160527 @1245		100		GM	DTW: 11' Screen: 29-39' First encountered groundwater:
35 - Cont Equif Drill Diami Logg	RACTOR Holt Services N PMENT Terra Sonic, track E . MTHD Sonic A ETER 8" B SER J Dahl REVIEWER C Bartlett	ORTHING ASTING NGLE EARING PRINTED	19985 13418 Vertic 07/08	3 58.780 396.090 cal 3/16	REMAR 05/27/2 monitor COORD SEE KEY S	KS: Sift 2016, ca ring we	e: Post office. We asing remained in t II 05/31/2016. SYSTEM: R SYMBOLS AND ABB	I Tag he bo	ID: BJ prehole	 X-186 05/28	. Comple -05/30, c	te drilling on onstructed

Ge	COSYNT consult	ants BOREHC	600	BORING START FINISH PROJEG LOCAT	DATE DATE DATE CT P ION Is CT NUM	MW04 5/27/2016 5/31/2016 hase 2 Hydroged ssaquah, Washin /IBER PNG0714	ologic gton 1	Elevat Invest	tion 7 igatior	SHE 73.30 FT	ET 2 OF 3		
DEPTH (ft)		MATERIAL DESCRIPTION	SYMBOLIC LOG	MELL LOG	W CONST MAT	ELL RUCTION ERIAL	ELEVATION (ft)	SAM SAMPLE NAME	PLES	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	wet, sandy GRA sorted, subround 6" gray, firm, 6" gray, firm, Gray (Gley 2 4/5 wet, SILT; trace fragments Gray (Gley 2 4/5 wet, sandy GRA cobbles Gray (Gley 2 4/5 wet, fine to coars graded 6" gray, firm, Gray (Gley 2 4/5 wet, SILT; some Gray (Gley 2 4/5 wet, SILT; some Gray (Gley 2 4/5 wet, sandy SILT Gray (Gley 2 4/5 wet, sandy SILT	WEL; well graded, poorly ded gravel wet, silt BBG dark greenish gray), firm, organics, trace wood BBG dark greenish gray), loose, VEL; well graded, trace BBG dark greenish gray), loose, se SAND; poorly sorted, well wet, silt BBG dark greenish gray), firm, organics BBG dark greenish gray), firm, e organics BBG dark greenish gray), firm, e silt			Halliburt Hole-plu bentonite (12-50lb	on g, 3/8" e chips bags)	- - - - - - - - - - - - - - - - - - -			100		ML GM SM ML SM SM-ML	33'
	Gray (Gley 2 4/5 medium SAND; Gray (Gley 2 4/5 wet, sandy SILT	BBG dark greenish gray), well sorted BBG dark greenish gray), firm,	• • • • • • • • • • • • • • • • •			_	- - 5 ⁻ -					SM SM-ML	
CONT EQUIF DRILL DIAMI LOGG	RACTOR Holt PMENT Terr MTHD Son ETER 8" ER J Dahl	t Services N ra Sonic, track E nic A REVIEWER C Bartlett	ORTHING ASTING NGLE EARING PRINTEE	i 1998 134 ² Vert 0 07/0	358.780 1896.090 ical 08/16	REMAR 05/27/2 monitor COORD SEE KEY S	KS: Sit 2016, ca ring we INATE	e: Post office. We asing remained in I 05/31/2016. SYSTEM: IR SYMBOLS AND ABB	ell Tag the bo	ID: BJ prehole	X-186. 05/28-	Comple 05/30, c	te drilling on onstructed

Ge	Geosyntec consultants GS FORM: CORE3 10/00 GS FORM: CORE3 10/00 CORE3 10/00 GS FORM: CORE3 10/00 CORE3 CORE CORE3 CORE3 COR							BORING MW04 SHEET 3 OF 3 START DATE 5/27/2016 Elevation 73.30 FT. MSL FINISH DATE 5/31/2016 PROJECT Phase 2 Hydrogeologic Investigation LOCATION Issaquah, Washington PROJECT NUMBER PNG0714						
DEPTH (ft)		MATERI/ DESCRIPT	AL ION	SYMBOLIC LOG	MELL LOG	W CONSTI MAT	ELL RUCTION ERIAL	ELEVATION (ft)	SAM SAMPLE NAME	PLES	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	Brown, medi well graded, Outwash) Brown, medi coarse SANI	ium dense, fine some rounded s ium dense, wet, D g, install monito	to coarse SAND; gravel (Advanced gravelly fine to			2" sch 44 0.01" slo screen	D PVC ttted well	- 0 			100		SP	
CONT EQUIP DRILL DIAME LOGG	RACTOR H MENT 7 MTHD 5 ETER 8 ER J Dahl	Holt Services Ferra Sonic, tr Sonic 3" REVIEWI	rack EA AN BE ER C Bartlett	DRTHING ASTING IGLE EARING PRINTED	19988 13418 Vertio 07/08	58.780 396.090 cal 3/16	REMAR 05/27/2 monito COORD SEE KEY S	KS: Sit 2016, ca ring we INATE	te: Post office. We asing remained in II 05/31/2016. SYSTEM: DR SYMBOLS AND ABB	ell Tag the bo	ID: BJ prehole	X-186 05/28	. Comple -05/30, c	ete drilling on constructed

Ge	Geosyntec consultants GS FORM: CORE3 10/00 GS FORM:						BORING START FINISH PROJEC LOCAT PROJEC	G DATE DATE CT P ION Is CT NUM	MW05 5/23/2016 5/23/2016 hase 2 Hydrogeo ssaquah, Washing MBER PNG0714	logic gton	Elevat Invest	ion igatio	SHE 72.05 F1 n	et 1 of 3 T. MSL
DEPTH (ft)		MATERIAL DESCRIPTION		SYMBOLIC LOG	MELL LOG	W CONSTI MAT	ELL RUCTION ERIAL	ELEVATION (ft)	SAMI SAMPLE NAME	PLES	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	Grass and To FILL (Light br GRAVEL; rou Dark gray, me GRAVEL/grav subangular gr Dark and light dark greenish fine SAND Gray (Gley 13 interbed with 1 Dark and light dark greenish fine SAND Very dark gray coarse SAND gravely SANU gravels Dark gray (Gle moist, SILT w fine sand inter Brown (10YR medium SANI angular sand Dark gray, me Medium brow medium dens rounded/subr trace silt, well quartzite, bas mound/subrour sit, well sorte basalt, and gr Dark gray, me 1 4/N) BRACTOR H	psoil own, loose, moist, s nd coarse gravel) adium dense, moist, relly SILT; trace fine avel, trace rounded gray mottled (Gley gray), medium den 3/N very dark gray) f trace firewood piece gray mottled (Gley gray), medium den 3/N very dark gray) f trace firewood piece gray mottled (Gley gray), medium den y (Gley 1 3/N), loose with some fine grav D, wet, well graded, ey 1 2.5/5GY greeni ith some clay; slight rbeds (<1") 4/2 dark grayish bro D; poorly graded, tra and fine subangular edium dense, wet, sa n (10YR 5/2 grayish e, wet, sandy GRAV bounded gravels, coarse sorted gravels, coarse sorted gravels of volcanic anodiorite dium dense, wet, si dium stiff, wet, sand Gravels, coarse s d gravels of volcanic anodiorite dium dense, wet, si dium stiff, wet, sand	ilty sandy silty sand, gravel 1 3/5GY very se, moist, silty ine SAND is 1 3/5GY very se, moist, silty ine SAND is 1 3/5GY very se, moist, silty is pround coarse sh black), plasticity, is plasticity, brown), loose, ace coarse gravels andy GRAVEL brown), /EL; rse sand, plcanics, brown), /EL; rse sand, plcanics, locanics, ity SAND to by SILT (Gley			Surface completi Morris monume 2" sch 4 well casi	on: 8" ent 0 PVC ng		MW05_10to20 _20160523 @1150		100 100	0 0 0 0 0 0 0 0 0	TOPSOIL FILL ML-GM SM SP SM SP ML SP GM GM GM SM-ML	easy drilling First groundwater: 10', rising DTW: 7.7' Screen: 10-20' 5' heave 5' heave
EQUIF DRILL DIAM LOGO	MENT TO MENT TO MTHD So ETER 8' ER J Dahl	erra Sonic, track onic REVIEWER C	NO EA: AN BE/ Bartlett P	STING STING GLE ARING RINTED	134 Ver 07/	1394.920 tical - 08/16		INATE	SYSTEM:	REVIAT	IONS	ט. ט. ד.	JV-103	

Ge	Geosyntec consultants GS FORM: CORE3 10/00 GS FORM: CORE3 CORE3						9600	BORING START FINISH PROJE LOCAT PROJE	G DATE DATE CT P TION Is	MW05 5/23/2016 5/23/2016 Phase 2 Hydrogeo ssaquah, Washing MBER PNG0714	ologic gton	Elevat Invest	ion igatio	SHE 72.05 F1 n	iet 2 of 3 T. MSL
	JRE3 10/00									SAM	PLES			uo)
DEPTH (ft)		D	MATERIAL		SYMBOLIC LOG	MELL LOG	CONS M/	WELL STRUCTION	ELEVATION (ft)	SAMPLE NAME	ТҮРЕ	% RECOVERY	PID READING (ppm)	USCS Classificati	COMMENTS
_	Brown (10Y medium SA	′R 4/3 ND; tr), medium dense, wet, fine race fine rounded gravels	to					_		\mathbb{N}			SP	
 40	Medium brc dense, wet, rounded grz Medium brc brown), me fine rounder Medium brc loose, wet, coarse sanc gravel	own/or coars avel, tr own/or dium o d grav own/or gravel d, coa	ange oxidation, medium ie sandy GRAVEL; fine ace coarse rounded gravel ange (7.5YR 4/6 strong Jense, wet, fine SAND; trac el ange (10YR 4/3 brown), ly SAND/sandy GRAVEL; rse rounded well graded	 ie /_			Hallibi Hole-p bentoi (14-50	urton Jlug, 3/8" lite chips Ib bags)	35 - - - - 30 -			100	0.1	GP SP GP	
 45 	Dark gray (medium stif fragments,	Gley 2 f, wet, slight	4/5B dark bluish gray), sandy SILT; trace wood sulfur odor		• • • • • • • • • • • • • • • • • • •				- - - 25 - -				0	SM-ML	
	Dark gray (medium stif	Gley 2 f, wet,	4/5B dark bluish gray), sandy SILT; trace organics						- - - 20 -			75	0	ML	
-	Dark gray (medium SA	Gley 2 f, wet,	, mediam dense, wet, ime ace gravel 4/5B dark bluish gray), sandy SILT	.u					-				0.1	SM-ML	
-	Brown (10Y SAND; well	'R 4/3 grade), medium dense, wet, silty d, poorly sorted, trace grav	el					- 15 ⁻ -				0	SM	
60 — — —	Brown (2.5)	ý 3/2 v ium S/	, very dark grayish brown), we AND; trace gravel, poorly	 et,					 10			100		SM	
 65	graded, wel Brown, mec oxidation at	ll sorte dium d conta	edense, wet, silty SAND;						-				0 0	SM	
	Brown (10Y sandy SILT Brown (10Y dense, wet, Gray (Gley sandy SII T	'R 5/4 , some 'R 5/4 silty c 2 4/5E	yellowish brown), stiff, wet, e wood fragments yellowish brown), medium coarse rounded GRAVELS 8 dark bluish gray), soft, wet	 「					5 - - -				0	SM-ML GP SM-ML GM	
CONT EQUIF	RACTOR	Holt : Terra	Services a Sonic, track	NOI EAS	MNYNYN RTHING STING	: 199 134	:] 057.680 1394.920	REMAR	 RKS: Si	l te: Salmon Run Pa	ark. W) /ell Tag	l ID: B	JX-183	
DRILL DIAMI LOGG	. MTHD ETER GER J Dahl	Sonio 8"	C REVIEWER C Bartlett	AN(BE/ P	GLE ARING RINTED	Ver 07	tical /08/16		DINATE	SYSTEM: DR SYMBOLS AND ABB	REVIAT	IONS			

Ge	Geosyntec consultants GS FORM: CORE3 10/00 GS FORM:						00	BORING START FINISH PROJEC LOCAT	G DATE DATE CT P ION Is	MW05 5/23/2016 5/23/2016 Phase 2 Hydroged ssaquah, Washin	ologic gton	Elevat Invest	tion igatio	SHE 72.05 F1 n	iet 3 of 3 r. MSL
	DRE3 10/00)				G		PROJE		WBER PNG0/14]
DEPTH (ft)		C	MATERIAL DESCRIPTION		SYMBOLIC LOG	MELL LOG	W CONSTI MAT	ell Ruction Erial	ELEVATION (ft)	SAM SAMPLE NAME	JES JAY	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	Brown (10' Imedium de Igraded, po Brown (10' trace cobb graded, rou Brown (10' well graded gravels Brown (10' medium S/ sorted, poc	YR 4/4 rnse, v orly sc YR 4/3 les, nc inded YR 4/3 d, pool YR 4/3 aND; t orly gra- ing, ins	dark yellowish brown), vet, silty sandy GRAVEL; w rrted	- I - J - J - J - J - To	෫෫෨෫෨෫෨෫෨෫෨෫෨෫෨෫෨෫෨෫෨෫෨෫෨෫෨෫෨෫෨෫෨෫෨෫෨෫		2" sch 4 0.01" slo screen	D PVC ttted well	- 0- - - -5- - - - -10- - - - - 15- - - - - - - - - - - - - -			75	0	SW	
CONT EQUIP DRILL DIAME LOGG	CONTRACTORHolt ServicesNORTHING199057.680EQUIPMENTTerra Sonic, trackEASTING1341394.920DRILL MTHDSonicANGLEVerticalDIAMETER8"BEARINGLOGGER J DahlREVIEWER C BartlettPRINTED07/08/16			57.680 394.920 cal 3/16	REMAR COORD SEE KEY S	KS: Si DINATE	te: Salmon Run Pa SYSTEM: DR SYMBOLS AND ABB	ark. W Reviat	ell Tag) ID: B	JX-183				

Geosyntec consultants GS FORM: CORE3 10/00 GS FORM: CORE3 CORE3 CO						BORING MW06 SHEET 1 OF 3 START DATE 10/5/2016 Elevation FT. MSL FINISH DATE 10/6/2016 PROJECT Phase 2 Hydrogeologic Investigation LOCATION Issaquah, Washington PROJECT NUMBER PNG0714							
	JRE3 10/00					(SAM	PLES			u	
DEPTH (ft)		MATERIAL DESCRIPTION	SYMBOLIC LOG	MELL LOG	W CONSTI MAT	ELL RUCTION ERIAL	ELEVATION (ft)	SAMPLE NAME	ТҮРЕ	% RECOVERY	PID READING (ppm)	USCS Classificatio	COMMENTS
-	FILL (Barkdust,	geotextile at 0.5') rown_loose_dry_silty_SAND	-		Surface		_		\mathbb{N}			FILL FILL	
-	topsoil)	iown, iouse, dry, sirry on vo,			completi Morris monume concrete	on: 8" ent, 2'x2' pad	-			100	0.3		
5 — -	Light olive brown SAND; trace silt	n (2.5Y 5/3), loose, dry, fine			Halliburt Hole-Plu bentonite (4-50lb t	on g, 3/8" e chips bags)	-			10	0.0	SM	Very soft
-							-			75			drilling, compressing samples, poor recovery
10 — 	Medium gray br fine sand, trace	own, soft, moist, SILT; some clay (slightly plastic)	+ + + + + + + + + + + + + + + + + + +				-			50	0.1	ML	
-	Dark bluish gray medium gravelly	/ (2 gley 4/5 PB), soft, moist, y SILT; less sand, more water, d brown trace claw, slightly					-			100		ML	
15 — 	plastic, oxidized	at lower contact					-			100	0.1		
-	Light brown gray large wood piec sulfur odor, trac	y, loose, moist, fine SAND; e at 19-19.5' (fresh), slight e silt, grading to coarse sand					-				0.3	SM	
20 —	with depth						-	CO1-MW06		100	1.0 0.1	GM	DTW: 20'
	some fine to coa sorted, occasior grades to coarse	arse sand, trace silt, poorly al coarse subrounded gravel, e gravel with depth					-	-20161005 -19.5-24.5 @1530		100	0.0		Screen: 19.5-24.5'
- 25	Brown (7.5YR 4	/4), loose, wet, sandy coarse					-			100	0.2	GW	
	GRAVEL; poorly subangular, trac	y sorted subrounded to be silt	20000000000000000000000000000000000000		2" sch 4		-			100	0.0	GW	Hit metal at 30'
	Dark yellow brow GRAVEL; some coarse subangu fine sand	wn (10YR 4/6), loose, wet, cobbles, coarse sand to lar to subrounded gravel, trace	4444		well cas	ng	-				0.2		Second hole on
			2000 2000 2000 2000 2000 2000 2000 200			_	-				0.3		metal at 32.5?
	RACTOR Cas	scade Drilling N				REMAR	KS: Si	te: Dogwood/New	oort. V	Vell Ta	g ID: E	3JH-349	
DRILL	. MTHD Sor		NGLE	Vei	rtical								
LOGG	ETER 6" ER C Bartlett	B REVIEWER C Bartlett	EARING PRINTED	11	 /07/16	COORD SEE KEY S	DINATE	SYSTEM: DR SYMBOLS AND ABE	REVIAT	IONS			

Ge	Geosyntec consultants GS FORM: CORE3 10/00 GS FORM: CORE3 10/00 GS FORM: CORE3 10/00 GS FORM: CORE3 10/00 GS FORM: COREAL C					BORING MW06 SHEET 2 OF 3 START DATE 10/5/2016 Elevation FT. MSL FINISH DATE 10/6/2016 PROJECT Phase 2 Hydrogeologic Investigation LOCATION Issaquah, Washington PROJECT NUMBER PNG0714					
DEPTH (ft)	MATERIAL DESCRIPTION	SYMBOLIC LOG	MELL LOG	WELL STRUCTION ATERIAL	ELEVATION (ft)	SAMF SAMFLE NAME	TYPE	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	- color change to dark brown (7.5YR 3/3), increased silt Dark brown, loose, wet, sandy GRAVEL; trace fine sand, trace subrounded cobbles Brown (7.5YR 4/3), loose, wet, silty sandy GRAVEL Dark brown, loose, wet, sandy GRAVEL; trace fine sand, trace subrounded cobbles Dark brown, loose, wet, coarse subrounded to subangular SAND; some fine gravel,				- - - - - - - - - - - - - - - - - - -	CO1-MW06 -20161006 -34.5-39.5 @1015		100 100 100	0.4 0.1 0.1 0.0 0.0 0.0	GW GP-GM GW GP GW SW	DTW: 30' Screen: 34.5-39.5'
	Brown, loose, gravelly SAND Dark brown, loose, wet, cobbly subangular GRAVEL; trace silt, trace fine sand Dark grayish brown (10YR 4/2), loose, wet, fine to medium SAND; trace fine gravel (no silt) Dark bluish gray (2 Gley 4/1 5PB), medium stiff, moist, fine sandy SILT; occasional fine rounded gravel		Hallit	ourton Grout Ib bags, ial of water)		CO1-MW06 -20161006 -51-56 @1235		100	0.0 0.0 0.0 0.0 0.0 0.0 0.0	SP GP-GM SP ML	DTW: 29' Screen: 51-56' 10' heave
CONT EQUIP DRILL DIAME LOGG	RACTOR Cascade Drilling NC PMENT Geoprobe 8140LS, track EA MTHD Sonic AN ETER 6" BE ER C Bartlett REVIEWER C Bartlett F	RTHING STING IGLE ARING PRINTED	Vertical 11/07/16	COORD SEE KEY S	KS: Si DINATE	te: Dogwood/Newp SYSTEM: DR SYMBOLS AND ABBI	ort. V	Vell Tag	g ID: E	3JH-349	



SONIC (PORTLAND) PNG0714 ISSAQUAH.GPJ EED DEFAULT GINT LIBRARY.GLB 11/7/16 I OG W/WFI I



Ge	Geosyntec consultants GS FORM: CORE3 10/00 GS FORM: CORE3 10/00 GS FORM: CORE3 10/00 GS FORM: CORE3 10/00 GS FORM: CORE3 10/00					BORING MW07 SHEET 2 OF 4 START DATE 10/7/2016 Elevation FT. MSL FINISH DATE 10/10/2016 PROJECT Phase 2 Hydrogeologic Investigation LOCATION Issaquah, Washington PROJECT NUMBER PNG0714					
DEPTH (ft)	MATERIAL DESCRIPTION	SYMBOLIC LOG	WELL LOG	WELL INSTRUCTION MATERIAL	ELEVATION (ft)	SAMF SAMFLE NAME	лтре Ттуре	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	Brown, loose, wet, silty fine to coarse subrounded GRAVEL to fine to coarse sand GRAVEL, with silt, occasional cobble	γ γ γ γ γ γ γ γ γ γ γ γ γ γ	<u>ЮПППППППППППППППППППППППППППППППППППП</u>	lliburton ik-Grout 50lb bags, 30 of water)		CO1-MW07 -20161007 -35-40 @1535		75		GM GM GM	DTW: 22' Screen: 35-40' 7' of heave, very soft, pushing out, poor recovery (depths recovered uncertain)
	Olive brown, very soft, wet, SILT Dark gray, soft, wet, clayey SILT; low to medium plasticity Dark gray (N4), medium stiff, moist, SILT; lo plasticity, trace very fine to fine sand				- - - - - - - - - - -	C01-MW07 -20161007 -65-70 @1800		25		ML ML	change to flapper bit very soft, fighting heave at 60', good recovery DTW: 52' Screen: 65-70'
CONT EQUIP DRILL DIAME LOGG	RACTOR Cascade Drilling PMENT Geoprobe 8140LS, track MTHD Sonic ETER 6" GER C Bartlett REVIEWER C Bartlett	NORTHING EASTING ANGLE BEARING PRINTEE	Vertical 0 11/07/16	COORD SEE KEY S	NINATE	te: Alder Court. Wo SYSTEM: DR SYMBOLS AND ABBI	ell Tag) ID: B	JH-348	}	



Ge	Geosyntec consultants GS FORM: CORE3 10/00 GS FORM:					BORING MW07 SHEET 4 OF 4 START DATE 10/7/2016 Elevation FT. MSL FINISH DATE 10/10/2016 PROJECT Phase 2 Hydrogeologic Investigation LOCATION Issaquah, Washington PROJECT NUMBER PNG0714							
DEPTH (ft)		MATERIAL DESCRIPTION	SYMBOLIC LOG	MELL LOG	W CONST MAT	ELL RUCTION ERIAL	ELEVATION (ft)	SAMPLE NAME	PLES	% RECOVERY	PID READING (ppm)	USCS Classification	COMMENTS
	End of borin	g, install monitoring well			2" sch 4(0.01" slo screen w 10/20 sil filter pac (4.5-50lb) PVC tted well vith ica sand k bags)				100			
CONT EQUIP DRILL DIAME LOGG	RACTOR (PMENT (MTHD (ETER (ER C Bartlet	Cascade Drilling Geoprobe 8140LS, track Sonic 5" tt REVIEWER C Bartlett	NORTHING EASTING ANGLE BEARING PRINTEI	Vertio 0 11/0	cal 7/16	REMAR COORD SEE KEY S	KS: Si INATE	te: Alder Court. W SYSTEM: DR SYMBOLS AND ABB	ell Taç REVIAT	g ID: B IONS	JH-34	8	

APPENDIX C ANALYTICAL DATA QUALITY REVIEW MEMOS, ANALYTICAL LABORATORY REPORTS



180A Market Place Boulevard Knoxville, TN 37922 PH 865.330.0037 www.geosyntec.com

Memorandum

- Date: 16 June 2016
- To: Bob Anderson
 - Cindy Bartlett
 - Samantha Fox
- From: Mary Tyler
- CC: J. Caprio
- Subject: Stage 2A Data Validation Level II Data Deliverables Eurofins Eaton Analytical Report Numbers 577272 Revised, 588864, 589670 and 591042

SITE: PNG0703

INTRODUCTION

This report summarizes the findings of the Stage 2A data validation of eight water samples, one field duplicate sample and four trip blank, collected February 19, 2016, May 4, 2016, May 10, 2016 and May 17, 2016, as part of the project PNG0703 sampling event. The analyses were performed at Eurofins Eaton Analytical, Monrovia, California. The samples were prepared and analyzed by the following analytical test:

• Perfluorinated alkyl acids (PFAAs) by EPA Method 537

EXECUTIVE SUMMARY

The samples were handled, prepared, and measured in the same manner under similar prescribed conditions.

Based on this Stage 2A data validation covering the quality control (QC) parameters listed below, the data are usable for meeting project objectives.

The organic data were reviewed based on the information and/or guidance provided in the US EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review, August 2014 (USEPA-540-R-014-002), as well as by the pertinent method referenced by the data package and technical and professional judgment.

The following samples were analyzed in the data set:

Laboratory ID	Client ID
201602240087	Well 4-021916
201602240088	Trip Blank - 021916
201605050734	Darigold-ABY249-050416
201605050735	Trip Blank - 050416
201605060067	Darigold-ABY249-050416-
	DUP
201605110272	SPWSD-VT2.1-051016
201605110273	SPWSD-VT2.2-051016

Laboratory ID	Client ID
201605110274	SPWSD-VT2.3-051016
201605110275	SPWSD-Well7.3-051016
201605110276	trip Blank - 051016
201605180426	COI-TW3-051716
201605180427	COI-MW1-051716
201605180428	TRIP BLANK-051716

The samples were received at the laboratory within the criteria of 0-6°C. No sample preservation issues were noted by the laboratory.

The laboratory noted that the samples reported in report 577272 were received partially frozen. This did not result in qualification of the data.

The chain of custody (COC) in report 577272 was not relinquished by the sampler.

Report 577272 was revised; the narrative indicated that the original reported perfluorooctanesulfonic acid (PFOS) result was mis-integrated; it did not include the branched isomers. The data for PFOS were re-integrated and revised results were reported in the revised laboratory report.

1.0 PERFLUORINATED ALKYL ACIDS

The samples were analyzed for PFAAs per EPA Method 537. The following compounds were reported:

- Perfluorobutanesulfonic acid, PFBS
- Perfluorodecanoic acid, PFDA
- Perfluorododecanoic acid, PFDoA
- Perfluoroheptanoic acid, PFHpA
- Perfluorohexanesulfonic acid, PFHxS
- Perfluorohexanoic acid, PFHxA
- Perfluorononanoic acid, PFNA
- Perfluorooctanesulfonic acid, PFOS
- Perfluorooctanoic acid, PFOA
- Perfluorotetradecanoic acid, PFTA
- Perfluorotridecanoic acid, PFTrDA
- Perfluoroundecanoic acid, PFUnA.

537 data DVR-Baseline

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The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Surrogates
- ✓ Trip Blank
- ✓ Equipment Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverables Review

1.1 <u>Overall Assessment</u>

The PFAA data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

1.2 <u>Holding Time</u>

The holding times for the PFAA analysis of a water sample are 14 days from collection to extraction and 28 days from extraction to analysis. The holding times were met for the sample analyses.

1.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Five method blanks were reported (batches 894999, 909576, 909665, 910527 and 911819). The PFAAs were not detected in the method blanks above the method reporting limits (MRLs).

537 data DVR-Baseline

1.4 <u>Matrix Spike/Matrix Spike Duplicate (MS/MSD)</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Batch MS/MSD pairs were reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

1.5 Laboratory Control Sample (LCS)

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Five LCS/LCS duplicate (LCSD) pairs were reported. The recovery and relative percent difference (RPD) results were within the method specified acceptance criteria.

1.6 <u>Surrogates</u>

Acceptable surrogate recoveries were reported for the sample analyses.

1.7 Equipment Blank

Equipment blanks were not collected with the sample sets.

1.8 <u>Trip Blank</u>

Four trip blanks, Trip Blank - 021916, Trip Blank – 050416, trip Blank – 051016 and TRIP BLANK-051716, accompanied the sample shipments. The PFAAs were not detected in the trip blanks above the MRLs.

1.9 <u>Field Duplicate</u>

One field duplicate sample, Darigold-ABY249-050416-DUP, was collected with the sample sets. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicate and the original sample, Darigold-ABY249-050416.

1.10 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

537 data DVR-Baseline

1.11 <u>Electronic Data Deliverables (EDD) Review</u>

Results and sample IDs in the EDDs were reviewed against the information provided by the associated level II reports at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II reports and the EDDs.

* * * * *

ATTACHMENT 1 DATA VALIDATION QUALIFIER DEFINITIONS AND INTERPRETATION KEY Assigned by Geosyntec's Data Validation Team

DATA QUALIFIER DEFINITIONS

- U The analyte was analyzed for, but was not detected above the reported sample quantitation limit.
- J The analyte was positively identified; the associated numerical value is the approximate concentration of the analyte in the sample.
- J+ The analyte was positively identified; however, the associated numerical value is likely to be higher than the concentration of the analyte in the sample due to positive bias of associated QC or calibration data or attributable to matrix interference.
- J- The analyte was positively identified; however, the associated numerical value is likely to be lower than the concentration of the analyte in the sample due to negative bias of associated QC or calibration data or attributable to matrix interference.
- UJ The analyte was not detected above the reported sample quantitation limit. However, the reported quantitation limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample.
- R The sample results are rejected due to serious deficiencies in the ability to analyze the sample and meet quality control criteria. The presence or absence of the analyte cannot be verified.
PNG0703 Data Validation 16 June 2016 Page 7

ATTACHMENT 2 DATA VALIDATION REASON CODES Assigned by Geosyntec's Data Validation Team

Valid Value	Description	
1	Preservation requirement not met	
2	Analysis holding time exceeded	
3	Blank contamination (i.e., method, trip, equipment, etc.)	
4	Matrix spike/matrix spike duplicate recovery or RPD outside limits	
5	LCS recovery outside limits	
6	Surrogate recovery outside limits	
7	Field Duplicate RPD exceeded	
8	Serial dilution percent difference exceeded	
9	Calibration criteria not met	
10	Linear range exceeded	
11	Internal standard criteria not met	
12	Lab duplicates RPD exceeded	
13	Other	

RPD-relative percent difference

537 data DVR-Baseline



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Memorandum

Subject:	Stage 2A Data Validation - Level 2 Data Deliverable – Eurofins Eaton Analytical Report Number 596940
CC:	J. Caprio
From:	Mary Tyler
	Cindy Bartlett
To:	Bob Anderson
Date:	30 August 2016

SITE: PNG0714

INTRODUCTION

This report summarizes the findings of the Stage 2A data validation of three water samples, one field duplicate sample and one trip blank, collected June 21, 2016, as part of the project PNG0714 sampling event. The analyses were performed at Eurofins Eaton Analytical, Monrovia, California. The samples were prepared and analyzed by the following analytical test:

• Perfluorinated alkyl acids (PFAAs) by EPA Method 537

EXECUTIVE SUMMARY

The samples were handled, prepared, and measured in the same manner under similar prescribed conditions.

Based on this Stage 2A data validation covering the quality control (QC) parameters listed below, the data are usable for meeting project objectives.

The organic data were reviewed based on the information and/or guidance provided in the US EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review, August 2014 (USEPA-540-R-014-002), as well as by the pertinent method referenced by the data package and technical and professional judgment.

The following samples were analyzed in the data set:

Laboratory ID	Client ID
201606220760	COI-MW03-20160621

Laboratory ID	Client ID	Laboratory ID	Client ID
201606220761	COI-MW03-20160621-	201606220763	COI-MW05-20160621
	DUP	201606220764	Trip Blank - 20160621
201606220762	COI-MW04-20160621		

The samples were received at the laboratory within the criteria of 0-6°C. No sample preservation issues were noted by the laboratory.

1.0 PERFLUORINATED ALKYL ACIDS

The samples were analyzed for PFAAs per EPA Method 537. The following compounds were reported:

- Perfluorobutanesulfonic acid, PFBS
- Perfluorodecanoic acid, PFDA
- Perfluorododecanoic acid, PFDoA
- Perfluoroheptanoic acid, PFHpA
- Perfluorohexanesulfonic acid, PFHxS
- Perfluorohexanoic acid, PFHxA
- Perfluorononanoic acid, PFNA
- Perfluorooctanesulfonic acid, PFOS
- Perfluorooctanoic acid, PFOA
- Perfluorotetradecanoic acid, PFTA
- Perfluorotridecanoic acid, PFTrDA
- Perfluoroundecanoic acid, PFUnA.

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Surrogates
- \checkmark Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverable Review

1.1 Overall Assessment

The PFAA data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

1.2 <u>Holding Time</u>

The holding times for the PFAA analysis of a water sample are 14 days from collection to extraction and 28 days from extraction to analysis. The holding times were met for the sample analyses.

1.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Three method blanks were reported (batches 918879, 918805 and 920999). The PFAAs were not detected in the method blanks above the method-specified one third (1/3) of the method reporting limits (MRLs).

1.4 <u>Matrix Spike/Matrix Spike Duplicate (MS/MSD)</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Batch MS/MSD pairs were reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

1.5 <u>Laboratory Control Sample (LCS)</u>

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LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Three LCS/LCS duplicate (LCSD) pairs were reported. The recovery and relative percent difference (RPD) results were within the method specified acceptance criteria.

1.6 <u>Surrogates</u>

Acceptable surrogate recoveries were reported for the sample analyses.

1.7 <u>Trip Blank</u>

One trip blank, Trip Blank - 20160621, accompanied the sample shipment. The PFAAs were not detected in the trip blank above the MRLs.

1.8 <u>Field Duplicate</u>

A field duplicate sample, COI-MW03-20160621-DUP, was collected with the sample set. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicate and the original sample, COI-MW03-20160621.

1.9 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

1.10 <u>Electronic Data Deliverable (EDD) Review</u>

Results and sample IDs in the EDD were reviewed against the information provided by the associated level IV report at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level IV report and the EDD.

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ATTACHMENT 1 DATA VALIDATION QUALIFIER DEFINITIONS AND INTERPRETATION KEY Assigned by Geosyntec's Data Validation Team

DATA QUALIFIER DEFINITIONS

- U The analyte was analyzed for, but was not detected above the reported sample quantitation limit.
- J The analyte was positively identified; the associated numerical value is the approximate concentration of the analyte in the sample.
- J+ The analyte was positively identified; however, the associated numerical value is likely to be higher than the concentration of the analyte in the sample due to positive bias of associated QC or calibration data or attributable to matrix interference.
- J- The analyte was positively identified; however, the associated numerical value is likely to be lower than the concentration of the analyte in the sample due to negative bias of associated QC or calibration data or attributable to matrix interference.
- UJ The analyte was not detected above the reported sample quantitation limit. However, the reported quantitation limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample.
- R The sample results are rejected due to serious deficiencies in the ability to analyze the sample and meet quality control criteria. The presence or absence of the analyte cannot be verified.

ATTACHMENT 2 DATA VALIDATION REASON CODES Assigned by Geosyntec's Data Validation Team

Valid Value	Description	
1	Preservation requirement not met	
2	Analysis holding time exceeded	
3	Blank contamination (i.e., method, trip, equipment, etc.)	
4	Matrix spike/matrix spike duplicate recovery or RPD outside limits	
5	LCS recovery outside limits	
6	Surrogate recovery outside limits	
7	Field Duplicate RPD exceeded	
8	Serial dilution percent difference exceeded	
9	Calibration criteria not met	
10	Linear range exceeded	
11	Internal standard criteria not met	
12	Lab duplicates RPD exceeded	
13	Other	

RPD-relative percent difference



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Memorandum

Subject:	Stage 2A Data Validation - Level 2 Data Deliverable – Eurofins Eaton Analytical Report Number 597948
CC:	J. Caprio
From:	Mary Tyler
	Cindy Bartlett
To:	Bob Anderson
Date:	30 August 2016

SITE: PNG0714

INTRODUCTION

This report summarizes the findings of the Stage 2A data validation of three water samples, one field duplicate sample and one trip blank, collected June 28, 2016, as part of the project PNG0714 sampling event. The analyses were performed at Eurofins Eaton Analytical, Monrovia, California. The samples were prepared and analyzed by the following analytical test:

• Perfluorinated alkyl acids (PFAAs) by EPA Method 537

EXECUTIVE SUMMARY

The samples were handled, prepared, and measured in the same manner under similar prescribed conditions.

Based on this Stage 2A data validation covering the quality control (QC) parameters listed below, the data are usable for meeting project objectives.

The organic data were reviewed based on the information and/or guidance provided in the US EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review, August 2014 (USEPA-540-R-014-002), as well as by the pertinent method referenced by the data package and technical and professional judgment.

The following samples were analyzed in the data set:

Laboratory ID	Client ID	Laboratory ID	Client ID
201606300054	COI-MW04-20160628	201606300055	COI-MW05-20160628

Laboratory ID	Client ID
201606300056	COI-MW03-20160628
201606300057	COI-MW03-20160628-
	DUP

Laboratory ID	Client ID
201606300058	Trip Blank-20160628

The samples were received at the laboratory within the criteria of 0-6°C. No sample preservation issues were noted by the laboratory.

1.0 PERFLUORINATED ALKYL ACIDS

The samples were analyzed for PFAAs per EPA Method 537. The following compounds were reported:

- Perfluorobutanesulfonic acid, PFBS
- Perfluorodecanoic acid, PFDA
- Perfluorododecanoic acid, PFDoA
- Perfluoroheptanoic acid, PFHpA
- Perfluorohexanesulfonic acid, PFHxS
- Perfluorohexanoic acid, PFHxA
- Perfluorononanoic acid, PFNA
- Perfluorooctanesulfonic acid, PFOS
- Perfluorooctanoic acid, PFOA
- Perfluorotetradecanoic acid, PFTA
- Perfluorotridecanoic acid, PFTrDA
- Perfluoroundecanoic acid, PFUnA.

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Surrogates
- \checkmark Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverable Review

1.1 Overall Assessment

The PFAA data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

1.2 <u>Holding Time</u>

The holding times for the PFAA analysis of a water sample are 14 days from collection to extraction and 28 days from extraction to analysis. The holding times were met for the sample analyses.

1.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). One method blank was reported (batch 920999). The PFAAs were not detected in the method blank above the method-specified one third (1/3) of the method reporting limits (MRLs).

1.4 <u>Matrix Spike/Matrix Spike Duplicate (MS/MSD)</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). A batch MS/MSD pair was reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

1.5 <u>Laboratory Control Sample (LCS)</u>

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). One LCS/LCS duplicate (LCSD) pair was reported. The recovery and relative percent difference (RPD) results were within the method specified acceptance criteria.

1.6 <u>Surrogates</u>

Acceptable surrogate recoveries were reported for the sample analyses.

1.7 <u>Trip Blank</u>

One trip blank, Trip Blank-20160628, accompanied the sample shipment. The PFAAs were not detected in the trip blanks above the MRLs.

1.8 <u>Field Duplicate</u>

A field duplicate sample, COI-MW03-20160628-DUP, was collected with the sample set. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicate and the original sample, COI-MW03-20160628.

1.9 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

1.10 <u>Electronic Data Deliverable (EDD) Review</u>

Results and sample IDs in the EDD were reviewed against the information provided by the associated level IV report at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level IV report and the EDD.

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ATTACHMENT 1 DATA VALIDATION QUALIFIER DEFINITIONS AND INTERPRETATION KEY Assigned by Geosyntec's Data Validation Team

DATA QUALIFIER DEFINITIONS

- U The analyte was analyzed for, but was not detected above the reported sample quantitation limit.
- J The analyte was positively identified; the associated numerical value is the approximate concentration of the analyte in the sample.
- J+ The analyte was positively identified; however, the associated numerical value is likely to be higher than the concentration of the analyte in the sample due to positive bias of associated QC or calibration data or attributable to matrix interference.
- J- The analyte was positively identified; however, the associated numerical value is likely to be lower than the concentration of the analyte in the sample due to negative bias of associated QC or calibration data or attributable to matrix interference.
- UJ The analyte was not detected above the reported sample quantitation limit. However, the reported quantitation limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample.
- R The sample results are rejected due to serious deficiencies in the ability to analyze the sample and meet quality control criteria. The presence or absence of the analyte cannot be verified.

ATTACHMENT 2 DATA VALIDATION REASON CODES Assigned by Geosyntec's Data Validation Team

Valid Value	Description	
1	Preservation requirement not met	
2	Analysis holding time exceeded	
3	Blank contamination (i.e., method, trip, equipment, etc.)	
4	Matrix spike/matrix spike duplicate recovery or RPD outside limits	
5	LCS recovery outside limits	
6	Surrogate recovery outside limits	
7	Field Duplicate RPD exceeded	
8	Serial dilution percent difference exceeded	
9	Calibration criteria not met	
10	Linear range exceeded	
11	Internal standard criteria not met	
12	Lab duplicates RPD exceeded	
13	Other	

RPD-relative percent difference



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Memorandum

Subject:	Stage 2A Data Validation - Level II Data Deliverables – Eurofins
CC:	J. Caprio
From:	Mary Tyler
	Cindy Bartlett
To:	Bob Anderson
Date:	30 August 2016

SITE: PNG0714

INTRODUCTION

This report summarizes the findings of the Stage 2A data validation of nine water samples, two field duplicate samples and two trip blanks, collected July 6, 2016 and July 13, 2016, as part of the project PNG0714 sampling event. The analyses were performed at Eurofins Eaton Analytical, Monrovia, California. The samples were prepared and analyzed by the following analytical tests:

Eaton Analytical Report Numbers 598922 and 600700

- Perfluorinated alkyl acids (PFAAs) by EPA Method 537
- Selected Metals (Calcium, Magnesium, Potassium, and Sodium) by EPA Method 200.7
- Anions by EPA Method 300.0

EXECUTIVE SUMMARY

The samples were handled, prepared, and measured in the same manner under similar prescribed conditions.

Based on this Stage 2A data validation covering the quality control (QC) parameters listed below, the data as qualified are usable for meeting project objectives. Qualified data should be used within the limitations of the qualification.

The data were reviewed based on the information and/or guidance provided in the US EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review, August 2014 (USEPA-540-R-014-002), the US EPA Contract Laboratory Program National Functional Guidelines for Inorganic Superfund Data Review, August 2014 (USEPA-540-

R-013-001), as well as by the pertinent methods referenced by the data package and technical and professional judgment.

Laboratory ID	Sample ID
201607070539	COI-MW04-20160716
201607070555	COI-Well 5-20160716
201607070556	COI-Well 50BS-20160716
201607070557	COI-MW05-20160716
201607070558	COI-MW03-20160716
201607070559	COI-MW03-20160716-Dup
201607070560	Trip Blank - 20160716
201607160575	COI-Well5-20160713

The following samples were analyzed in the data set:

Laboratory ID	Sample ID
201607160577	COI-MW04-20160713
201607160578	COI-MW05-20160713
201607160579	COI-MW03-20160713
201607160580	COI-MW03-20160713 DUP
201607160581	TRIP BLANK-20160713

The samples were received at the laboratory within the criteria of 0-6°C. No sample preservation issues were noted by the laboratory.

The chain of custody for the samples reported in laboratory report number 600700 indicated metals and anion analyses for trip blank TRIP BLANK-20160713. Metals and anions were not reported for this sample.

1.0 PERFLUORINATED ALKYL ACIDS

The samples were analyzed for PFAAs per EPA Method 537. The following compounds were reported:

- Perfluorobutanesulfonic acid, PFBS
- Perfluorodecanoic acid, PFDA
- Perfluorododecanoic acid, PFDoA
- Perfluoroheptanoic acid, PFHpA
- Perfluorohexanesulfonic acid, PFHxS
- Perfluorohexanoic acid, PFHxA
- Perfluorononanoic acid, PFNA
- Perfluorooctanesulfonic acid, PFOS
- Perfluorooctanoic acid, PFOA
- Perfluorotetradecanoic acid, PFTA
- Perfluorotridecanoic acid, PFTrDA
- Perfluoroundecanoic acid, PFUnA.

598922 600700 DVR R1

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The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- \otimes Surrogates
- ✓ Trip Blank
- \otimes Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverables Review

1.1 Overall Assessment

The PFAA data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

1.2 <u>Holding Time</u>

The holding times for the PFAA analysis of a water sample are 14 days from collection to extraction and 28 days from extraction to analysis. The holding times were met for the sample analyses.

1.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Three method blanks were reported (batches 921762, 923973 and 923636). The PFAAs were not detected in the method blanks above the method-specified one third (1/3) of the method reporting limits (MRLs).

598922 600700 DVR R1

1.4 <u>Matrix Spike/Matrix Spike Duplicate (MS/MSD)</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Batch MS/MSD pairs were reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

1.5 Laboratory Control Sample (LCS)

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Three LCS/LCS duplicate (LCSD) pairs were reported. The recovery and relative percent difference (RPD) results were within the method specified acceptance criteria.

1.6 <u>Surrogates</u>

Acceptable surrogate recoveries were reported for the sample analyses, with the following exception.

The recovery of 13C-PFDA in sample COI-MW03-20160716-Dup was low and outside the method specified acceptance criteria. Therefore, based on professional and technical judgment, the nondetect result of PFDA in sample COI-MW03-20160716-Dup was J qualified as estimated.

Sample ID	Compound	Laboratory Result (µg/L)	Laboratory Flag	Validation Result (µg/L)	Validation Qualifier*	Reason Code**
COI-MW03- 20160716-Dup	Perfluorodecanoic acid	0.0025	US7	0.0025	UJ	6

µg/L-micrograms per liter

U-not detected at or above the MRL

*Validation qualifiers are defined in Attachment 1 at the end of this report

** Reason codes are defined in Attachment 2 at the end of this report

1.7 <u>Trip Blank</u>

Two trip blanks, Trip Blank – 20160716 and TRIP BLANK-20160713, were collected with the sample sets. The PFAAs were not detected in the trip blanks above the MRLs.

1.8 <u>Field Duplicate</u>

Two field duplicate samples, COI-MW03-060716-DUP and COI-MW03-20160713 DUP, were collected with the sample sets. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated

598922 600700 DVR R1

between the field duplicates and the original samples, COI-MW03-060716 and COI-MW03-20160713, with the following exception.

The RPD for perfluorooctanoic acid was greater than 30% and the concentrations in the field duplicate pair were greater than two times the MRL. Therefore, the concentrations of perfluorooctanoic acid in the field duplicate pair were J qualified as estimated.

Sample ID	Compound	Laboratory	Laboratory	RPD	Validation	Validation	Reason
_	_	Result	Flag		Result	Qualifier	Code
		(µg/L)			(µg/L)		
COI-MW03-	Perfluorobutanesulfonic	0.025	NA	4	NA	NA	NA
20160716	acid						
COI-MW03-	Perfluorobutanesulfonic	0.026	S7		NA	NA	NA
20160716-Dup	acid						
COI-MW03-	Perfluoroheptanoic acid	0.0063	NA	2	NA	NA	NA
20160716							
COI-MW03-	Perfluoroheptanoic acid	0.0062	S7		NA	NA	NA
20160716-Dup							
COI-MW03-	Perfluorohexanesulfonic	0.062	NA	0	NA	NA	NA
20160716	acid						
COI-MW03-	Perfluorohexanesulfonic	0.062	S7		NA	NA	NA
20160716-Dup	acid						
COI-MW03-	Perfluorohexanoic acid	0.015	NA	7	NA	NA	NA
20160716							
COI-MW03-	Perfluorohexanoic acid	0.014	S7		NA	NA	NA
20160716-Dup							
COI-MW03-	Perfluorononanoic acid	0.0061	NA	9	NA	NA	NA
20160716							
COI-MW03-	Perfluorononanoic acid	0.0056	S7		NA	NA	NA
20160716-Dup							
COI-MW03-	Perfluorooctanesulfonic	0.10	NA	0	NA	NA	NA
20160716	acid						
COI-MW03-	Perfluorooctanesulfonic	0.10	NA		NA	NA	NA
20160716-Dup	acid						
COI-MW03-	Perfluorooctanoic acid	0.0051	NA	33	0.0051	J	7
20160716							
COI-MW03-	Perfluorooctanoic acid	0.0071	S7		0.0071	J	7
20160716-Dup							
COI-MW03-	The other PFAAs	ND	NA	0	NA	NA	NA
20160716							
COI-MW03-	The other PFAAs	ND	NA		NA	NA	NA
20160716-Dup							
COI-MW03-	Perfluorobutanesulfonic	0.031	NA	7	NA	NA	NA
20160713	acid						
COI-MW03-	Perfluorobutanesulfonic	0.029	NA		NA	NA	NA
20160713 DUP	acid						

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Sample ID	Compound	Laboratory Result	Laboratory Flag	RPD	Validation Result	Validation Qualifier	Reason Code
		(µg/L)	8		(µg/L)		
COI-MW03-	Perfluoroheptanoic acid	0.0071	NA	10	NA	NA	NA
20160713							
COI-MW03-	Perfluoroheptanoic acid	0.0064	NA		NA	NA	NA
20160713 DUP							
COI-MW03-	Perfluorohexanesulfonic	0.075	NA	7	NA	NA	NA
20160713	acid						
COI-MW03-	Perfluorohexanesulfonic	0.070	NA		NA	NA	NA
20160713 DUP	acid						
COI-MW03-	Perfluorohexanoic acid	0.016	NA	13	NA	NA	NA
20160713							
COI-MW03-	Perfluorohexanoic acid	0.014	NA		NA	NA	NA
20160713 DUP							
COI-MW03-	Perfluorononanoic acid	0.0063	NA	14	NA	NA	NA
20160713							
COI-MW03-	Perfluorononanoic acid	0.0055	NA		NA	NA	NA
20160713 DUP							
COI-MW03-	Perfluorooctanesulfonic	0.088	NA	13	NA	NA	NA
20160713	acid						
COI-MW03-	Perfluorooctanesulfonic	0.10	NA		NA	NA	NA
20160713 DUP	acid						
COI-MW03-	Perfluorooctanoic acid	0.0053	NA	12	NA	NA	NA
20160713 DUP							
COI-MW03-	Perfluorooctanoic acid	0.0060	NA		NA	NA	NA
20160713							
COI-MW03-	The other PFAAs	ND	NA	0	NA	NA	NA
20160713 DUP							
COI-MW03-	The other PFAAs	ND	NA		NA	NA	NA
20160713							

µg/L-micrograms per liter

NA-not applicable

S7-laborartory flag indicating the surrogate recovery was below laboratory and method acceptance limits; unable to confirm matrix effect

ND-not detected at or above the MRL

1.9 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

1.10 Electronic Data Deliverables (EDD) Review

Results and sample IDs in the EDDs were reviewed against the information provided by the associated level II reports at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II reports and the EDDs.

2.0 SELECTED METALS

The samples were analyzed for selected metals per EPA Method 200.7.

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverables Review

2.1 Overall Assessment

The metals data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

2.2 <u>Holding Time</u>

The holding time for the metals analysis of a water sample is 180 days from collection to analysis. The holding times were met for the sample analyses.

2.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Four method blanks were reported (batches 921963, 922648, 924020 and 924992). The metals were not detected in the method blanks above the MRLs.

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2.4 <u>Matrix Spike/Matrix Spike Duplicate</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Batch MS/MSD pairs were reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

2.5 <u>Laboratory Control Sample</u>

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Four LCS/LCSD pairs were reported. The recovery and RPD results were within the method specified acceptance criteria.

2.6 <u>Trip Blank</u>

The trip blanks were not analyzed for metals.

2.7 <u>Field Duplicate</u>

Two field duplicate samples, COI-MW03-060716-DUP and COI-MW03-20160713 DUP, were collected with the sample sets. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicates and the original samples, COI-MW03-060716 and COI-MW03-20160713.

2.8 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

2.9 <u>Electronic Data Deliverables Review</u>

Results and sample IDs in the EDDs were reviewed against the information provided by the associated level II reports at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II reports and the EDDs.

3.0 ANIONS

The samples were analyzed for anions by EPA method 300.0.

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Final Review: JK Caprio 09/02/16

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The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ⊗ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverables Review

3.1 Overall Assessment

The anion data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

3.2 <u>Holding Time</u>

The holding times for water samples are listed below.

Analysis	Holding Time
Nitrate as N, Nitrate as NO ₃ , and Nitrite as N and total nitrate, nitrate-	48 hours from collection to analysis
N by calculation by EPA Method 300.0	
Chloride and Sulfate by EPA Method 300.0	28 days from collection to analysis

The holding times were met for the sample analyses, with the following exception. Sample COI-Well5-20160713 was analyzed fifty-two hours after collection. Therefore, the nondetect results of nitrate and nitrite in this sample were UJ qualified as estimated less than the MRLs.

Sample ID	Compound	Laboratory Result (mg/L)	Laboratory Flag	Validation Result (mg/L)	Validation Qualifier	Reason Code
COI-Well5- 20160713	Nitrate (as N)	0.10	UH3	0.10	UJ	2
COI-Well5- 20160713	Nitrate (as NO ₃)	0.44	UH3	0.44	UJ	2
COI-Well5- 20160713	Nitrite (as N)	0.050	UH3	0.050	UJ	2
COI-Well5- 20160713	Nitrate and Nitrite (as N)	0.10	UH3	0.10	UJ	2

mg/L-milligrams per liter

U-not detected at or above the MRL

H3-laboratory flag indicating the sample was received and/or analysis requested past holding time

3.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Four method blanks were reported (batches 921569, 921570, 923178 and 923179). The anions were not detected in the method blanks above the MRLs.

3.4 <u>Matrix Spike/Matrix Spike Duplicate</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Sample specific MS/MSD pairs were reported for the anions using sample COI-MW04-20160716. The recovery and RPD results were within the laboratory specified acceptance criteria.

Batch MS/MSD pairs were also reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

3.5 Laboratory Control Sample

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Four LCS/LCSD pairs were reported. The recovery and RPD results were within the method specified acceptance criteria.

3.6 <u>Trip Blank</u>

The trip blanks were not analyzed for metals.

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Final Review: JK Caprio 09/02/16

3.7 <u>Field Duplicate</u>

Two field duplicate samples, COI-MW03-060716-DUP and COI-MW03-20160713 DUP, were collected with the sample sets. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicates and the original samples, COI-MW03-060716 and COI-MW03-20160713.

3.8 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

3.9 <u>Electronic Data Deliverables Review</u>

Results and sample IDs in the EDDs were reviewed against the information provided by the associated level II reports at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II reports and the EDDs.

* * * * *

ATTACHMENT 1 DATA VALIDATION QUALIFIER DEFINITIONS AND INTERPRETATION KEY Assigned by Geosyntec's Data Validation Team

DATA QUALIFIER DEFINITIONS

- U The analyte was analyzed for, but was not detected above the reported sample quantitation limit.
- J The analyte was positively identified; the associated numerical value is the approximate concentration of the analyte in the sample.
- J+ The analyte was positively identified; however, the associated numerical value is likely to be higher than the concentration of the analyte in the sample due to positive bias of associated QC or calibration data or attributable to matrix interference.
- J- The analyte was positively identified; however, the associated numerical value is likely to be lower than the concentration of the analyte in the sample due to negative bias of associated QC or calibration data or attributable to matrix interference.
- UJ The analyte was not detected above the reported sample quantitation limit. However, the reported quantitation limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample.
- R The sample results are rejected due to serious deficiencies in the ability to analyze the sample and meet quality control criteria. The presence or absence of the analyte cannot be verified.

ATTACHMENT 2 DATA VALIDATION REASON CODES Assigned by Geosyntec's Data Validation Team

Valid Value	Description
1	Preservation requirement not met
2	Analysis holding time exceeded
3	Blank contamination (i.e., method, trip, equipment, etc.)
4	Matrix spike/matrix spike duplicate recovery or RPD outside limits
5	LCS recovery outside limits
6	Surrogate recovery outside limits
7	Field Duplicate RPD exceeded
8	Serial dilution percent difference exceeded
9	Calibration criteria not met
10	Linear range exceeded
11	Internal standard criteria not met
12	Lab duplicates RPD exceeded
13	Other

RPD-relative percent difference

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Memorandum

Date:	07 September 2016

To: Bob Anderson Cindy Bartlett

From: Mary Tyler

CC: J. Caprio

Subject: Stage 2A Data Validation - Level II Data Deliverables – Eurofins Eaton Analytical Report Number 601877

SITE: PNG0714

INTRODUCTION

This report summarizes the findings of the Stage 2A data validation of three water samples, one field duplicate sample and one trip blank, collected July 20, 2016, as part of the project PNG0714 sampling event. The analyses were performed at Eurofins Eaton Analytical, Monrovia, California. The samples were prepared and analyzed by the following analytical tests:

- Perfluorinated alkyl acids (PFAAs) by EPA Method 537
- Selected Metals (Calcium, Magnesium, Potassium, and Sodium) by EPA Method 200.7
- Anions by EPA Method 300.0

EXECUTIVE SUMMARY

The samples were handled, prepared, and measured in the same manner under similar prescribed conditions.

Based on this Stage 2A data validation covering the quality control (QC) parameters listed below, the data are usable for meeting project objectives.

The data were reviewed based on the information and/or guidance provided in the US EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review, August 2014 (USEPA-540-R-014-002), the US EPA Contract Laboratory Program National Functional Guidelines for Inorganic Superfund Data Review, August 2014 (USEPA-540-R-013-001), as well as by the pertinent methods referenced by the data package and technical and professional judgment.

The following samples were analyzed in the data set:

Laboratory ID	Sample ID
201607220374	COI-MW04-20160720
201607220376	COI-MW05-20160720
201607220377	COI-MW03-20160720
201607220378	COI-MW03-20160720-DUP

Laboratory ID	Sample ID
201607220379	TRIP BLANK - 20160720

The samples were received at the laboratory within the criteria of 0-6°C. No sample preservation issues were noted by the laboratory.

The following issues were noted with the chain of custody (COC); these did not result in qualification of the data.

- The COC indicated that the anion analyses were on hold; anions were reported for the samples.
- The COC indicated metals and anion analyses for trip blank TRIP BLANK-20160720. Metals and anions were not reported for this sample. In addition, no collection time was listed on the COC for the trip blank. The laboratory assigned a collection time of 0800.

1.0 PERFLUORINATED ALKYL ACIDS

The samples were analyzed for PFAAs per EPA Method 537. The following compounds were reported:

- Perfluorobutanesulfonic acid, PFBS
- Perfluorodecanoic acid, PFDA
- Perfluorododecanoic acid, PFDoA
- Perfluoroheptanoic acid, PFHpA
- Perfluorohexanesulfonic acid, PFHxS
- Perfluorohexanoic acid, PFHxA
- Perfluorononanoic acid, PFNA
- Perfluorooctanesulfonic acid, PFOS
- Perfluorooctanoic acid, PFOA
- Perfluorotetradecanoic acid, PFTA
- Perfluorotridecanoic acid, PFTrDA
- Perfluoroundecanoic acid, PFUnA.

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues

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were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Surrogates
- ✓ Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverables Review

1.1 Overall Assessment

The PFAA data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

1.2 <u>Holding Time</u>

The holding times for the PFAA analysis of a water sample are 14 days from collection to extraction and 28 days from extraction to analysis. The holding times were met for the sample analyses.

1.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Two method blanks were reported (batches 925241 and 925697). The PFAAs were not detected in the method blanks above the method-specified one third (1/3) of the method reporting limits (MRLs).

1.4 Matrix Spike/Matrix Spike Duplicate (MS/MSD)

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Batch MS/MSD pairs were reported. Since these are

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batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

1.5 <u>Laboratory Control Sample (LCS)</u>

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Two LCS/LCS duplicate (LCSD) pairs were reported. The recovery and relative percent difference (RPD) results were within the method specified acceptance criteria.

1.6 <u>Surrogates</u>

Acceptable surrogate recoveries were reported for the sample analyses.

1.7 <u>Trip Blank</u>

A trip blank, TRIP BLANK-20160720, was collected with the sample set. The PFAAs were not detected in the trip blank above the MRLs.

1.8 <u>Field Duplicate</u>

A field duplicate sample, COI-MW03-20160720-DUP, was collected with the sample set. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicate and the original sample, COI-MW03-20160720.

1.9 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

1.10 Electronic Data Deliverable (EDD) Review

Results and sample IDs in the EDD were reviewed against the information provided by the associated level II report at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II report and the EDD.

2.0 SELECTED METALS

The samples were analyzed for selected metals per EPA Method 200.7.

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Final Review: JK Caprio 09/08/16

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- \checkmark Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverables Review

2.1 Overall Assessment

The metals data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

2.2 <u>Holding Time</u>

The holding time for the metals analysis of a water sample is 180 days from collection to analysis. The holding times were met for the sample analyses.

2.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Two method blanks were reported (batches 925591 and 925849). The metals were not detected in the method blanks above the MRLs.

2.4 <u>Matrix Spike/Matrix Spike Duplicate</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Batch MS/MSD pairs were reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

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2.5 <u>Laboratory Control Sample</u>

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Two LCS/LCSD pairs were reported. The recovery and RPD results were within the method specified acceptance criteria.

2.6 <u>Trip Blank</u>

The trip blank was not analyzed for metals.

2.7 <u>Field Duplicate</u>

A field duplicate sample, COI-MW03-20160720-DUP, was collected with the sample set. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicate and the original sample, COI-MW03-20160720.

2.8 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

2.9 <u>Electronic Data Deliverable Review</u>

Results and sample IDs in the EDD were reviewed against the information provided by the associated level II report at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II report and the EDD.

3.0 ANIONS

The samples were analyzed for anions by EPA method 300.0.

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate

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Final Review: JK Caprio 09/08/16

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- ✓ Laboratory Control Sample
- ✓ Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverables Review

3.1 Overall Assessment

The anion data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

3.2 <u>Holding Time</u>

The holding times for water samples are listed below. The holding times were met for the sample analyses. It was noted that the nitrate and nitrite analysis of sample COI-MW04-20160720 was performed 18 minutes outside the 48 hour holding time. Since it was analyzed within the hour of the 48 hours, no qualifications were applied to the data, based on professional and technical judgment.

Analysis	Holding Time
Nitrate as N, Nitrate as NO ₃ , and Nitrite as N and total nitrate, nitrate-	48 hours from collection to analysis
N by calculation by EPA Method 300.0	
Chloride and Sulfate by EPA Method 300.0	28 days from collection to analysis

3.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Two method blanks were reported (batches 924873 and 924874). The anions were not detected in the method blanks above the MRLs.

3.4 <u>Matrix Spike/Matrix Spike Duplicate</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Batch MS/MSD pairs were reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

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3.5 <u>Laboratory Control Sample</u>

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Two LCS/LCSD pairs were reported. The recovery and RPD results were within the method specified acceptance criteria.

3.6 <u>Trip Blank</u>

The trip blank was not analyzed for anions.

3.7 <u>Field Duplicate</u>

A field duplicate sample, COI-MW03-20160720-DUP, was collected with the sample set. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicate and the original sample, COI-MW03-20160720.

3.8 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

3.9 <u>Electronic Data Deliverable Review</u>

Results and sample IDs in the EDD were reviewed against the information provided by the associated level II report at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II report and the EDD.

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ATTACHMENT 1 DATA VALIDATION QUALIFIER DEFINITIONS AND INTERPRETATION KEY Assigned by Geosyntec's Data Validation Team

DATA QUALIFIER DEFINITIONS

- U The analyte was analyzed for, but was not detected above the reported sample quantitation limit.
- J The analyte was positively identified; the associated numerical value is the approximate concentration of the analyte in the sample.
- J+ The analyte was positively identified; however, the associated numerical value is likely to be higher than the concentration of the analyte in the sample due to positive bias of associated QC or calibration data or attributable to matrix interference.
- J- The analyte was positively identified; however, the associated numerical value is likely to be lower than the concentration of the analyte in the sample due to negative bias of associated QC or calibration data or attributable to matrix interference.
- UJ The analyte was not detected above the reported sample quantitation limit. However, the reported quantitation limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample.
- R The sample results are rejected due to serious deficiencies in the ability to analyze the sample and meet quality control criteria. The presence or absence of the analyte cannot be verified.

ATTACHMENT 2 DATA VALIDATION REASON CODES Assigned by Geosyntec's Data Validation Team

Valid Value	Description
1	Preservation requirement not met
2	Analysis holding time exceeded
3	Blank contamination (i.e., method, trip, equipment, etc.)
4	Matrix spike/matrix spike duplicate recovery or RPD outside limits
5	LCS recovery outside limits
6	Surrogate recovery outside limits
7	Field Duplicate RPD exceeded
8	Serial dilution percent difference exceeded
9	Calibration criteria not met
10	Linear range exceeded
11	Internal standard criteria not met
12	Lab duplicates RPD exceeded
13	Other

RPD-relative percent difference

601877 DVR


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Memorandum

Subject:	Stage 2A Data Validation - Level 2 Data Deliverable – Eurofins Lancaster Laboratories Analytical Report Number 601992
CC:	J. Caprio
From:	Mary Tyler
	Cindy Bartlett
To:	Bob Anderson
Date:	07 September 2016

SITE: PNG0714

INTRODUCTION

This report summarizes the findings of the Stage 2A data validation of three soil samples and one field duplicate sample, collected July 22, 2016, as part of the project PNG0714 sampling event. The analyses were performed at Eurofins Lancaster Laboratories, Lancaster, Pennsylvania. The samples were prepared and analyzed by the following analytical test:

• Perfluorinated alkyl acids (PFAAs) by EPA Method 537, revision 1.1 Mod

EXECUTIVE SUMMARY

The samples were handled, prepared, and measured in the same manner under similar prescribed conditions.

Based on this Stage 2A data validation covering the quality control (QC) parameters listed below, the data as qualified are usable for meeting project objectives. Qualified data should be used within the limitations of the qualifications.

The organic data were reviewed based on the information and/or guidance provided in the US EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review, August 2014 (USEPA-540-R-014-002), as well as by the pertinent method referenced by the data package and technical and professional judgment.

The following samples were analyzed in the data set:

Laboratory ID	Client ID
201607230255	COI-STTA01-20160722
201607230256	COI-STTA02-20160722
201607230257	COI-STTA02-20160722-
	DUP

Laboratory ID	Client ID
201607230258	COI-STSP01-20160722

The samples were received at the laboratory within the criteria of 0-6°C. No sample preservation issues were noted by the laboratory.

1.0 PERFLUORINATED ALKYL ACIDS

The samples were analyzed for PFAAs per EPA Method 537, revision 1.1 Mod. The following compounds were reported:

- Perfluorobutanesulfonic acid, PFBS
- Perfluorodecanoic acid, PFDA
- Perfluorododecanoic acid, PFDoA
- Perfluoroheptanoic acid, PFHpA
- Perfluorohexanesulfonic acid, PFHxS
- Perfluorohexanoic acid, PFHxA
- Perfluorononanoic acid, PFNA
- Perfluorooctanesulfonic acid, PFOS
- Perfluorooctanoic acid, PFOA
- Perfluorotetradecanoic acid, PFTA
- Perfluorotridecanoic acid, PFTrDA
- Perfluoroundecanoic acid, PFUnA

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Surrogates
- ✓ Trip Blank
- ⊗ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverable Review

601992 DVR

engineers | scientists | innovators

Final Review:

1.1 Overall Assessment

The PFAA data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

1.2 <u>Holding Time</u>

The method lists the holding times for the PFAA analysis of a water sample. Additional information from the laboratory indicated that the holding times for soils are 28 days from collection to extraction and 28 days from extraction to analysis. The laboratory specified holding times were met for the soil sample analyses.

1.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). One method blank was reported (batch 16210006). The PFAAs were not detected in the method blank above the limits of quantitation [LOQs, also called the reporting limits (RLs)].

1.4 <u>Matrix Spike (MS)</u>

A sample set specific MS was reported, using sample COI-STTA01-20160722. The recovery results were within the laboratory specified acceptance criteria, with the following exception.

The recovery of perfluorooctanesulfonic acid was high and outside the laboratory specified acceptance criteria. Due to the difference between the sample and spike concentrations and professional and technical judgment, no qualifications were applied to the data.

It was noted that the sample results for COI-STTA01-20160722 were reported on a dry weight basis and the MS results were reported on a wet weight basis. This did not result in qualification of the data.

1.5 Laboratory Control Sample (LCS)

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). One LCS/LCS duplicate (LCSD) pair was reported. The recovery and relative percent difference (RPD) results were within the method specified acceptance criteria.

1.6 <u>Surrogates</u>

Acceptable surrogate recoveries were reported for the sample analyses.

1.7 <u>Trip Blank</u>

A trip blank did not accompany the sample shipment.

1.8 <u>Field Duplicate</u>

A field duplicate sample, COI-STTA02-20160722-DUP, was collected with the sample set. Acceptable precision (RPD \leq 50%) was demonstrated between the field duplicate and the original sample, COI-STTA02-20160722, with the following exception.

The RPD for perfluoroundecanoic acid was greater than 50%. Therefore, based on professional and technical judgment, the concentrations of perfluoroundecanoic acid in the field duplicate pair were J qualified as estimated.

Sample ID	Compound	Laboratory	Laboratory	RPD	Validation	Validation	Reason Codo**
		(ng/g)	гад		(ng/g)	Quanneation	Coue
COI- STTA02- 20160722	PFBS	4.5	NA	18	NA	NA	NA
COI- STTA02- 20160722- DUP	PFBS	5.4	NA		NA	NA	NA
COI- STTA02- 20160722	PFDA	3.9	NA	14	NA	NA	NA
COI- STTA02- 20160722- DUP	PFDA	4.5	NA		NA	NA	NA
COI- STTA02- 20160722	PFHpA	2.1	NA	13	NA	NA	NA
COI- STTA02-	PFHpA	2.4	NA		NA	NA	NA

601992 DVR

Final Review: JK Caprio 09/08/16

Sample ID	Compound	Laboratory Concentration	Laboratory Flag	RPD	Validation Concentration	Validation Oualification*	Reason Code**
		(ng/g)			(ng/g)		
20160722-							
DUP							
COI-	PFHxS	25	NA	15	NA	NA	NA
STTA02-							
20160722	DEIL	20	NIA		NA	NA	NA
STTA02	РГПХЗ	29	NA		NA	INA	INA
20160722-							
DUP							
COI-	PFHxA	15	NA	24	NA	NA	NA
STTA02-							
20160722							
COI-	PFHxA	19	NA		NA	NA	NA
STTA02-							
20160722-							
DUP	DENIA	22	27.4	26	274		27.4
COI-	PFNA	33	NA	26	NA	NA	NA
STTA02- 20160722							
20100722	ΡΕΝΛ	13	ΝA	_	ΝΔ	NΔ	ΝA
STTA02-	TIMA	45	INA		INA	INA	INA
20160722-							
DUP							
COI-	PFOS	180	NA	33	NA	NA	NA
STTA02-							
20160722							
COI-	PFOS	250	NA		NA	NA	NA
STTA02-							
20160722-							
DUP	DEOA	4.2	NIA	10	NIA	NIA	NA
STTA02	PTOA	4.5	NA	19	NA	INA	INA
20160722							
COI-	PFOA	5.2	NA		NA	NA	NA
STTA02-		0.2				1.1.1	1.1.1
20160722-							
DUP							
COI-	PFTrDA	2.7	NA	11	NA	NA	NA
STTA02-							
20160722							
COI-	PFTrDA	3	NA		NA	NA	NA
STTA02-							
20100722-							
COI-	PFUnA	36	NA	55	36	T	7
STTA02-	TIUIA	50		55	50	5	/
20160722							
COI-	PFUnA	63	NA	1	63	J	7
STTA02-							
20160722-							
DUP							

601992 DVR

Sample ID	Compound	Laboratory Concentration (ng/g)	Laboratory Flag	RPD	Validation Concentration (ng/g)	Validation Qualification*	Reason Code**
COI- STTA02- 20160722	The other PFAAs	ND	NA	0	NA	NA	NA
COI- STTA02- 20160722- DUP	The other PFAAs	ND	NA		NA	NA	NA

ng/g -nanograms per gram

NA-not applicable

ND-not detected

* Validation qualifiers are defined in Attachment 1 at the end of this report

**Reason codes are defined in Attachment 2 at the end of this report

1.9 <u>Sensitivity</u>

The samples were reported to the LOQs. No elevated nondetect results were reported for the samples. It was noted that the nondetect results for sample COI-STSP01-20160722 were reported as ND (\leq); the nondetect results in the other samples were reported as ND. Based on information from the laboratory, this was due to the data entered in the laboratory information system (LIMS) by the analyst; this did not result in qualifications of the data.

1.10 Electronic Data Deliverable (EDD) Review

Results and sample IDs in the EDD were reviewed against the information provided by the associated level IV report at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level IV report and the EDD.

* * * * *

ATTACHMENT 1 DATA VALIDATION QUALIFIER DEFINITIONS AND INTERPRETATION KEY Assigned by Geosyntec's Data Validation Team

DATA QUALIFIER DEFINITIONS

- U The analyte was analyzed for, but was not detected above the reported sample quantitation limit.
- J The analyte was positively identified; the associated numerical value is the approximate concentration of the analyte in the sample.
- J+ The analyte was positively identified; however, the associated numerical value is likely to be higher than the concentration of the analyte in the sample due to positive bias of associated QC or calibration data or attributable to matrix interference.
- J- The analyte was positively identified; however, the associated numerical value is likely to be lower than the concentration of the analyte in the sample due to negative bias of associated QC or calibration data or attributable to matrix interference.
- UJ The analyte was not detected above the reported sample quantitation limit. However, the reported quantitation limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample.
- R The sample results are rejected due to serious deficiencies in the ability to analyze the sample and meet quality control criteria. The presence or absence of the analyte cannot be verified.

ATTACHMENT 2 DATA VALIDATION REASON CODES Assigned by Geosyntec's Data Validation Team

Valid Value	Description
1	Preservation requirement not met
2	Analysis holding time exceeded
3	Blank contamination (i.e., method, trip, equipment, etc.)
4	Matrix spike/matrix spike duplicate recovery or RPD outside limits
5	LCS recovery outside limits
6	Surrogate recovery outside limits
7	Field Duplicate RPD exceeded
8	Serial dilution percent difference exceeded
9	Calibration criteria not met
10	Linear range exceeded
11	Internal standard criteria not met
12	Lab duplicates RPD exceeded
13	Other

RPD-relative percent difference



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Memorandum

Subject:	Stage 2A Data Validation - Level 2 Data Deliverable – Eurofins Eaton Analytical Report Number 602965
CC:	J. Caprio
From:	Mary Tyler
	Cindy Bartlett
To:	Bob Anderson
Date:	03 September 2016

SITE: PNG0714

INTRODUCTION

This report summarizes the findings of the Stage 2A data validation of four water samples, one field duplicate sample and one trip blank, collected July 27-28, 2016, as part of the project PNG0714 sampling event. The analyses were performed at Eurofins Eaton Analytical, Monrovia, California. The samples were prepared and analyzed by the following analytical test:

• Perfluorinated alkyl acids (PFAAs) by EPA Method 537

EXECUTIVE SUMMARY

The samples were handled, prepared, and measured in the same manner under similar prescribed conditions.

Based on this Stage 2A data validation covering the quality control (QC) parameters listed below, the data are usable for meeting project objectives.

The organic data were reviewed based on the information and/or guidance provided in the US EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review, August 2014 (USEPA-540-R-014-002), as well as by the pertinent method referenced by the data package and technical and professional judgment.

The following samples were analyzed in the data set:

Laboratory ID	Client ID	Laboratory ID	Client ID
201607290381	COI-MW04-20160727	201607290382	COI-Well5-20160727

Laboratory ID	Client ID
201607290383	Trip Blank - 20160727
201607290389	COI-MW03-20160728
201607290390	COI-MW03-20160728-
	DUP

Laboratory ID	Client ID
201607290391	COI-MW05-20160728

The samples were received at the laboratory within the criteria of 0-6°C. No sample preservation issues were noted by the laboratory.

The following issues were noted with the chain of custody (COC) form; no qualifications were applied to the data:

- Analyses and matrices were noted for samples COI-MW04-20160727 and COI-MW05-20160728 on the COC. PFAAs were reported the other water samples listed on the COC.
- There was no time of collection listed for the trip blank on the COC; the laboratory assigned a collection time of 0800.

1.0 PERFLUORINATED ALKYL ACIDS

The samples were analyzed for PFAAs per EPA Method 537. The following compounds were reported:

- Perfluorobutanesulfonic acid, PFBS
- Perfluorodecanoic acid, PFDA
- Perfluorododecanoic acid, PFDoA
- Perfluoroheptanoic acid, PFHpA
- Perfluorohexanesulfonic acid, PFHxS
- Perfluorohexanoic acid, PFHxA
- Perfluorononanoic acid, PFNA
- Perfluorooctanesulfonic acid, PFOS
- Perfluorooctanoic acid, PFOA
- Perfluorotetradecanoic acid, PFTA
- Perfluorotridecanoic acid, PFTrDA
- Perfluoroundecanoic acid, PFUnA

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Surrogates
- ✓ Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverable Review

1.1 Overall Assessment

The PFAA data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

1.2 <u>Holding Time</u>

The holding times for the PFAA analysis of a water sample are 14 days from collection to extraction and 28 days from extraction to analysis. The holding times were met for the sample analyses.

1.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Two method blanks were reported (batches 926352 and 926827). The PFAAs were not detected in the method blanks above the method-specified one third (1/3) of the method reporting limits (MRLs).

1.4 <u>Matrix Spike/Matrix Spike Duplicate (MS/MSD)</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Batch MS/MSD pairs were reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

1.5 Laboratory Control Sample (LCS)

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Two LCS/LCS duplicate (LCSD) pairs were reported. The recovery and relative percent difference (RPD) results were within the method specified acceptance criteria.

1.6 <u>Surrogates</u>

Acceptable surrogate recoveries were reported for the sample analyses.

1.7 <u>Trip Blank</u>

One trip blank, Trip Blank - 20160727, accompanied the sample shipment. The PFAAs were not detected in the trip blank above the MRLs.

1.8 <u>Field Duplicate</u>

A field duplicate sample, COI-MW03-20160728-DUP, was collected with the sample set. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicate and the original sample, COI-MW03-20160728.

1.9 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

1.10 Electronic Data Deliverable (EDD) Review

Results and sample IDs in the EDD were reviewed against the information provided by the associated level IV report at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level IV report and the EDD.

* * * * *

ATTACHMENT 1 DATA VALIDATION QUALIFIER DEFINITIONS AND INTERPRETATION KEY Assigned by Geosyntec's Data Validation Team

DATA QUALIFIER DEFINITIONS

- U The analyte was analyzed for, but was not detected above the reported sample quantitation limit.
- J The analyte was positively identified; the associated numerical value is the approximate concentration of the analyte in the sample.
- J+ The analyte was positively identified; however, the associated numerical value is likely to be higher than the concentration of the analyte in the sample due to positive bias of associated QC or calibration data or attributable to matrix interference.
- J- The analyte was positively identified; however, the associated numerical value is likely to be lower than the concentration of the analyte in the sample due to negative bias of associated QC or calibration data or attributable to matrix interference.
- UJ The analyte was not detected above the reported sample quantitation limit. However, the reported quantitation limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample.
- R The sample results are rejected due to serious deficiencies in the ability to analyze the sample and meet quality control criteria. The presence or absence of the analyte cannot be verified.

ATTACHMENT 2 DATA VALIDATION REASON CODES Assigned by Geosyntec's Data Validation Team

Valid Value	Description
1	Preservation requirement not met
2	Analysis holding time exceeded
3	Blank contamination (i.e., method, trip, equipment, etc.)
4	Matrix spike/matrix spike duplicate recovery or RPD outside limits
5	LCS recovery outside limits
6	Surrogate recovery outside limits
7	Field Duplicate RPD exceeded
8	Serial dilution percent difference exceeded
9	Calibration criteria not met
10	Linear range exceeded
11	Internal standard criteria not met
12	Lab duplicates RPD exceeded
13	Other

RPD-relative percent difference



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Memorandum

- Date: 24 June 2016
- To: Bob Anderson
 - Cindy Bartlett

Samantha Fox

- From: Mary Tyler
- CC: J. Caprio
- Subject: Stage 2A Data Validation Level II Data Deliverables Eurofins Eaton Analytical Report Numbers 592302, 592671, 593456, and 594492

SITE: PNG0714

INTRODUCTION

This report summarizes the findings of the Stage 2A data validation of twelve water samples, one field duplicate sample, one equipment blank, and four trip blanks, collected May 23 – June 7, 2016, as part of the project PNG0714 sampling event. The analyses were performed at Eurofins Eaton Analytical, Monrovia, California. The samples were prepared and analyzed by the following analytical tests:

- Perfluorinated alkyl acids (PFAAs) by EPA Method 537
- Selected Volatile Organic Compounds (VOCs) [Benzene, Toluene, Ethylbenzene, and Xylenes (BTEX) and methyl tert butyl ether (MTBE)] by EPA Method 524.2
- Selected Metals (Calcium, Magnesium, Potassium, and Sodium) by EPA Method 200.7
- Anion by EPA Method 300.0
- Alkalinity by Standard Method 2320B
- Specific Conductance by Standard Method 2510B
- pH by Standard Method 4500-HB

EXECUTIVE SUMMARY

The samples were handled, prepared, and measured in the same manner under similar prescribed conditions.

Based on this Stage 2A data validation covering the quality control (QC) parameters listed below, the data are usable for meeting project objectives.

The organic data were reviewed based on the information and/or guidance provided in the US EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review, August 2014 (USEPA-540-R-014-002), as well as by the pertinent method referenced by the data package and technical and professional judgment.

The inorganic data were reviewed based on the information and/or guidance provided in the US EPA Contract Laboratory Program National Functional Guidelines for Inorganic Superfund Data Review, August 2014 (USEPA-540-R-013-001), as well as by the pertinent method referenced by the data package and technical and professional judgment.

The following samples were analyzed in the data set:

Laboratory ID	Sample ID
201605260223	MW05_10 to 20_20160523
201605260227	MW03_10 to 20_20160524
201605260228	TRIP BLANK_20160524
201605260229	MW03_40 to 50_20160524
201605280242	MW01_30 to 40_20160526
201605280243	MW01_Rinseate_20160526
201605280244	MW01_55 to 65_20160526
201605280245	MW04_29 to 39_20160527
201605280246	trip blank_20160527
201606020681	MW02_40to50_20160531

Laboratory ID	Sample ID
201606020682	Trip Blank_20160601
201606080558	COI-MW01-060716
201606080560	COI-MW02-060716
201606080561	COI-MW03-060716
201606080562	COI-MW03-060716-DUP
201606080563	COI-MW04-060716
201606080564	COI-MW05-060716
201606080568	Trip Blank-060716

The samples were received at the laboratory within the criteria of 0-6°C. No sample preservation issues were noted by the laboratory.

The relinquished date and time were not listed on the chain of custody (COC) in report 592302.

1.0 PERFLUORINATED ALKYL ACIDS

The samples were analyzed for PFAAs per EPA Method 537. The following compounds were reported:

- Perfluorobutanesulfonic acid, PFBS
- Perfluorodecanoic acid, PFDA
- Perfluorododecanoic acid, PFDoA
- Perfluoroheptanoic acid, PFHpA

- Perfluorohexanesulfonic acid, PFHxS
- Perfluorohexanoic acid, PFHxA
- Perfluorononanoic acid, PFNA
- Perfluorooctanesulfonic acid, PFOS
- Perfluorooctanoic acid, PFOA
- Perfluorotetradecanoic acid, PFTA
- Perfluorotridecanoic acid, PFTrDA
- Perfluoroundecanoic acid, PFUnA.

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Surrogates
- ✓ Equipment Blank
- \otimes Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverables Review

1.1 Overall Assessment

The PFAA data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

1.2 <u>Holding Time</u>

The holding times for the PFAA analysis of a water sample are 14 days from collection to extraction and 28 days from extraction to analysis. The holding times were met for the sample analyses.

1.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Nine method blanks were reported (batches 914151, 914427, 915130, 915652, 915136, 916189, 916843, 916753, and 917227). The PFAAs were not detected in the method blanks above the method reporting limits (MRLs).

1.4 <u>Matrix Spike/Matrix Spike Duplicate (MS/MSD)</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Batch MS/MSD pairs were reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

It was noted that some MS/MSD recovery calculations appeared incorrect based on the data presented. The laboratory has not provided information explaining the inconsistencies. Since these are batch QC the reports were validated based on the pertinent associated QC samples and no qualifications were applied to the data.

1.5 <u>Laboratory Control Sample (LCS)</u>

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Nine LCS/LCS duplicate (LCSD) pairs were reported. The recovery and relative percent difference (RPD) results were within the method specified acceptance criteria.

1.6 <u>Surrogates</u>

Acceptable surrogate recoveries were reported for the sample analyses, with the following exception.

The recovery of 13C-PFDA in sample MW01_Rinseate_20160526 was high and outside the laboratory specified acceptance criteria. Since the remaining three surrogates were within laboratory limits, no qualifications were applied to the data based on professional judgment.

1.7 Equipment Blank

One equipment blank, MW01_Rinseate_20160526, was collected with the sample set. The PFAAs were not detected in the equipment blank above the MRLs.

1.8 <u>Trip Blank</u>

Four trip blanks, TRIP BLANK_20160524, trip blank_20160527, Trip Blank_20160601, and Trip Blank-060716, accompanied the sample shipments. The PFAAs were not detected in the trip blanks above the MRLs, with the following exceptions.

Perfluoroheptanoic acid (0.0099 μ g/L), perfluorohexanesulfonic acid (0.023 $\mu g/L$), perfluorohexanoic acid (0.021)μg/L), perfluorononanoic acid (0.0085) $\mu g/L$), perfluorooctanesulfonic acid (0.059 µg/L), and perfluorooctanoic acid (0.011 µg/L) were detected at concentrations greater than the MRLs in trip blank 20160527. Since perfluoroheptanoic acid, perfluorohexanoic acid, perfluorononanoic acid, and perfluorooctanoic acid were not detected in the associated samples, no qualifications were applied to these compounds. However, the perfluorohexanesulfonic acid and perfluorooctanesulfonic acid concentrations greater than the MRLs and less than the trip blank contamination in the associated sample were U qualified as not detected at the reported concentrations.

Sample ID	Compound	Laboratory Result (µg/L)	Laboratory Flag	Validation Result (µg/L)	Validation Qualifier*	Reason Code**
MW04_29 to	Perfluorohexanesulfonic	0.0084	NA	0.0084	U	3
39_20160527	acid					
MW04_29 to	Perfluorooctanesulfonic	0.0028	NA	0.0028	U	3
39 20160527	acid					

µg/L-micrograms per liter

NA-not applicable

*Validation qualifiers are defined in Attachment 1 at the end of this report

** Reason codes are defined in Attachment 2 at the end of this report

1.9 <u>Field Duplicate</u>

One field duplicate sample, COI-MW03-060716-DUP, was collected with the sample sets. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicate and the original sample, COI-MW03-060716.

1.10 Sensitivity

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

1.11 <u>Electronic Data Deliverables (EDD) Review</u>

Results and sample IDs in the EDDs were reviewed against the information provided by the associated level II reports at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II reports and the EDDs.

2.0 SELECTED VOLATILE ORGANIC COMPOUNDS

The samples were analyzed for selected VOCs per EPA Method 524.2.

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- \otimes Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Surrogates
- ✓ Trip Blank
- ✓ Equipment Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverables Review

2.1 <u>Overall Assessment</u>

The VOC data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

2.2 <u>Holding Time</u>

The holding time for the VOC analysis of a water sample is 14 days from collection to analysis. The holding times were met for the sample analyses, with the following exception.

Sample COI-MW03-060716-DUP was Q3 flagged by the laboratory indicating the sample was not properly preserved. The laboratory indicated the sample pH was greater than 2; therefore, the sample was considered unpreserved. The holding time for the VOC analysis of an unpreserved water sample is 7 days from collection to analysis. The sample was analyzed outside the holding time. Therefore, the nondetect results in sample COI-MW03-060716-DUP were UJ qualified as estimated less than the MRL based on professional judgment.

Sample ID	Compound	Laboratory Result (µg/L)	Laboratory Flag	Validation Result (µg/L)	Validation Qualifier	Reason Code
COI-MW03-060716- DUP	Benzene	0.50	UQ3	0.50	UJ	2
COI-MW03-060716- DUP	Ethylbenzene	0.50	UQ3	0.50	UJ	2
COI-MW03-060716- DUP	m&p-Xylenes	0.50	UQ3	0.50	UJ	2
COI-MW03-060716- DUP	Methyl tert-Butyl Ether	0.50	UQ3	0.50	UJ	2
COI-MW03-060716- DUP	o-Xylene	0.50	UQ3	0.50	UJ	2
COI-MW03-060716- DUP	Toluene	0.50	UQ3	0.50	UJ	2
COI-MW03-060716- DUP	Xylenes, Total	0.50	UQ3	0.50	UJ	2

µg/L-micrograms per liter

U-not detected at or above the MRL

Q3-laboratory flag indicating sample was not properly preserved

2.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). One method blank was reported (batch 917124). The VOCs were not detected in the method blank above the MRLs.

2.4 <u>Matrix Spike/Matrix Spike Duplicate</u>

MS/MSD pairs were not reported.

2.5 <u>Laboratory Control Sample</u>

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). One LCS/LCSD pair was reported. The recovery and RPD results were within the method specified acceptance criteria.

Issaquah DVR Stage 2A June 2016

2.6 <u>Surrogates</u>

Acceptable surrogate recoveries were reported for the sample analyses.

2.7 Equipment Blank

The equipment blank was not analyzed for VOCs.

2.8 <u>Trip Blank</u>

One trip blank, Trip Blank-060716, accompanied the sample shipments and was analyzed for VOCs. The VOCs were not detected in the trip blank above the MRLs.

2.9 <u>Field Duplicate</u>

One field duplicate sample, COI-MW03-060716-DUP, was collected with the sample sets. Acceptable precision (RPD \leq 30%) was demonstrated between the field duplicate and the original sample, COI-MW03-060716. The RPDs were 0%.

2.10 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

2.11 <u>Electronic Data Deliverables Review</u>

Results and sample IDs in the EDDs were reviewed against the information provided by the associated level II reports at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II reports and the EDDs.

3.0 SELECTED METALS

The samples were analyzed for selected metals per EPA Method 200.7.

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time

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- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Equipment Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverables Review

3.1 <u>Overall Assessment</u>

The metals data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

3.2 <u>Holding Time</u>

The holding time for the metals analysis of a water sample is 180 days from collection to analysis. The holding times were met for the sample analyses.

3.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). One method blank was reported (batch 916682). The metals were not detected in the method blank above the MRLs.

3.4 <u>Matrix Spike/Matrix Spike Duplicate</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Batch MS/MSD pairs were reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

3.5 <u>Laboratory Control Sample</u>

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). One LCS/LCSD pair was reported. The recovery and RPD results were within the method specified acceptance criteria.

3.6 Equipment Blank

The equipment blank was not analyzed for metals.

3.7 <u>Field Duplicate</u>

One field duplicate sample, COI-MW03-060716-DUP, was collected with the sample set. Acceptable precision (RPD \leq 30%) was demonstrated between the field duplicate and the original sample, COI-MW03-060716.

3.8 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

3.9 <u>Electronic Data Deliverables Review</u>

Results and sample IDs in the EDDs were reviewed against the information provided by the associated level II reports at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II reports and the EDDs.

4.0 WET CHEMISTRY PARAMETERS

The samples were analyzed for anions by EPA method 300.0, alkalinity by standard method 2320B, specific conductance by standard method 2510B, and pH by Standard Method 4500-HB.

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Equipment Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverables Review

4.1 Overall Assessment

The wet chemistry data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

4.2 <u>Holding Time</u>

The holding times for water samples are listed below. The holding times were met for the sample analyses.

Analysis	Holding Time
Nitrate as N, Nitrate as NO ³ , and Nitrite as N by EPA Method 300.0	48 hours from collection to analysis
Chloride and Sulfate by EPA Method 300.0	28 days from collection to analysis
Alkalinity by Standard Method 2320B	14 days from collection to analysis
Specific Conductivity by Standard Method 2510B	28 days from collection to analysis
pH by Standard Method 4500-HB	As soon as possible

pH does not have a specific holding time listed in method 4500-HB; the method indicates the samples should be analyzed as soon as possible after sample collection since pH changes occur within 15 minutes of sampling. Based on professional and technical judgment, as soon as possible should be within one day of receipt by the laboratory at most. Therefore, since the water pH analyses were performed within one day of laboratory receipt, no qualifications were applied to the water pH data.

4.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Four method blanks were reported (batches 915726, 915736, 915950, and 915960). The wet chemistry parameters were not detected in the method blanks above the MRLs.

4.4 <u>Matrix Spike/Matrix Spike Duplicate</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Sample specific MS/MSD pairs were reported for

nitrate as N, nitrite as N, chloride, and sulfate using sample COI-MW02-060716. The recovery and RPD results were within the laboratory specified acceptance criteria.

Batch MS/MSD pairs were also reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

4.5 <u>Laboratory Control Sample</u>

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Five LCS/LCSD pairs were reported. The recovery and RPD results were within the method specified acceptance criteria.

4.6 <u>Laboratory Duplicate</u>

Batch laboratory duplicates were reported for pH and specific conductivity. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

4.7 <u>Equipment Blank</u>

The equipment blank was not analyzed for metals.

4.8 <u>Field Duplicate</u>

One field duplicate sample, COI-MW03-060716-DUP, was collected with the sample sets. Acceptable precision (RPD \leq 30%) was demonstrated between the field duplicate and the original sample, COI-MW03-060716.

4.9 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

4.10 <u>Electronic Data Deliverables Review</u>

Results and sample IDs in the EDDs were reviewed against the information provided by the associated level II reports at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II reports and the EDDs.

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ATTACHMENT 1 DATA VALIDATION QUALIFIER DEFINITIONS AND INTERPRETATION KEY Assigned by Geosyntec's Data Validation Team

DATA QUALIFIER DEFINITIONS

- U The analyte was analyzed for, but was not detected above the reported sample quantitation limit.
- J The analyte was positively identified; the associated numerical value is the approximate concentration of the analyte in the sample.
- J+ The analyte was positively identified; however, the associated numerical value is likely to be higher than the concentration of the analyte in the sample due to positive bias of associated QC or calibration data or attributable to matrix interference.
- J- The analyte was positively identified; however, the associated numerical value is likely to be lower than the concentration of the analyte in the sample due to negative bias of associated QC or calibration data or attributable to matrix interference.
- UJ The analyte was not detected above the reported sample quantitation limit. However, the reported quantitation limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample.
- R The sample results are rejected due to serious deficiencies in the ability to analyze the sample and meet quality control criteria. The presence or absence of the analyte cannot be verified.

ATTACHMENT 2 DATA VALIDATION REASON CODES Assigned by Geosyntec's Data Validation Team

Valid Value	Description
1	Preservation requirement not met
2	Analysis holding time exceeded
3	Blank contamination (i.e., method, trip, equipment, etc.)
4	Matrix spike/matrix spike duplicate recovery or RPD outside limits
5	LCS recovery outside limits
6	Surrogate recovery outside limits
7	Field Duplicate RPD exceeded
8	Serial dilution percent difference exceeded
9	Calibration criteria not met
10	Linear range exceeded
11	Internal standard criteria not met
12	Lab duplicates RPD exceeded
13	Other

RPD-relative percent difference

Issaquah DVR Stage 2A June 2016



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Memorandum

Subject:	Stage 2A Data Validation - Level II Data Deliverable – Eurofins Eaton Analytical Report Number 613606
CC:	J. Caprio
From:	Mary Tyler
	Cindy Bartlett
To:	Bob Anderson
Date:	07 November 2016

SITE: PNG0714

INTRODUCTION

This report summarizes the findings of the Stage 2A data validation of two water samples and one trip blank, collected September 27, 2016, as part of the project PNG0714 sampling event. The analyses were performed at Eurofins Eaton Analytical, Monrovia, California. The samples were prepared and analyzed by the following analytical test:

• Perfluorinated alkyl acids (PFAAs) by EPA Method 537

EXECUTIVE SUMMARY

The samples were handled, prepared, and measured in the same manner under similar prescribed conditions.

Based on this Stage 2A data validation covering the quality control (QC) parameters listed below, the data are usable for meeting project objectives.

The data were reviewed based on the information and/or guidance provided in the US EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review, August 2014 (USEPA-540-R-014-002), as well as by the pertinent method referenced by the data package and technical and professional judgment.

The following samples were analyzed in the data set:

Laboratory ID	Sample ID	Laboratory ID	Sample ID
201609280826	COI-SW-A-092716	201609280851	COI-SW-B-092716

Laboratory ID	Sample ID
201609280852	TripBlank-092716

The samples were received at the laboratory within the criteria of 0-6°C. No sample preservation issues were noted by the laboratory.

1.0 PERFLUORINATED ALKYL ACIDS

The samples were analyzed for PFAAs per EPA Method 537. The following compounds were reported:

- Perfluorobutanesulfonic acid, PFBS
- Perfluorodecanoic acid, PFDA
- Perfluorododecanoic acid, PFDoA
- Perfluoroheptanoic acid, PFHpA
- Perfluorohexanesulfonic acid, PFHxS
- Perfluorohexanoic acid, PFHxA
- Perfluorononanoic acid, PFNA
- Perfluorooctanesulfonic acid, PFOS
- Perfluorooctanoic acid, PFOA
- Perfluorotetradecanoic acid, PFTA
- Perfluorotridecanoic acid, PFTrDA
- Perfluoroundecanoic acid, PFUnA.

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Surrogates
- ✓ Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverable Review

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1.1 Overall Assessment

The PFAA data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

1.2 <u>Holding Time</u>

The holding times for the PFAA analysis of a water sample are 14 days from collection to extraction and 28 days from extraction to analysis. The holding times were met for the sample analyses.

1.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). One method blank was reported (batch 940060). The PFAAs were not detected in the method blank above the method-specified one third (1/3) of the method reporting limits (MRLs).

1.4 <u>Matrix Spike/Matrix Spike Duplicate (MS/MSD)</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). One batch MS/MSD pair was reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

1.5 <u>Laboratory Control Sample (LCS)</u>

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). One LCS/LCS duplicate (LCSD) pair was reported. The recovery and relative percent difference (RPD) results were within the method specified acceptance criteria.

1.6 <u>Surrogates</u>

Acceptable surrogate recoveries were reported for the sample analyses.

1.7 <u>Trip Blank</u>

A trip blank, TripBlank-092716, was collected with the sample set. The PFAAs were not detected in the trip blank above the MRLs.

1.8 <u>Field Duplicate</u>

A field duplicate sample was not collected with the sample set.

1.9 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

1.10 Electronic Data Deliverable (EDD) Review

Results and sample IDs in the EDD were reviewed against the information provided by the associated level II report at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II report and the EDD.

* * * * *

ATTACHMENT 1 DATA VALIDATION QUALIFIER DEFINITIONS AND INTERPRETATION KEY Assigned by Geosyntec's Data Validation Team

DATA QUALIFIER DEFINITIONS

- U The analyte was analyzed for, but was not detected above the reported sample quantitation limit.
- J The analyte was positively identified; the associated numerical value is the approximate concentration of the analyte in the sample.
- J+ The analyte was positively identified; however, the associated numerical value is likely to be higher than the concentration of the analyte in the sample due to positive bias of associated QC or calibration data or attributable to matrix interference.
- J- The analyte was positively identified; however, the associated numerical value is likely to be lower than the concentration of the analyte in the sample due to negative bias of associated QC or calibration data or attributable to matrix interference.
- UJ The analyte was not detected above the reported sample quantitation limit. However, the reported quantitation limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample.
- R The sample results are rejected due to serious deficiencies in the ability to analyze the sample and meet quality control criteria. The presence or absence of the analyte cannot be verified.

ATTACHMENT 2 DATA VALIDATION REASON CODES Assigned by Geosyntec's Data Validation Team

Valid Value	Description
1	Preservation requirement not met
2	Analysis holding time exceeded
3	Blank contamination (i.e., method, trip, equipment, etc.)
4	Matrix spike/matrix spike duplicate recovery or RPD outside limits
5	LCS recovery outside limits
6	Surrogate recovery outside limits
7	Field Duplicate RPD exceeded
8	Serial dilution percent difference exceeded
9	Calibration criteria not met
10	Linear range exceeded
11	Internal standard criteria not met
12	Lab duplicates RPD exceeded
13	Other

RPD-relative percent difference



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Memorandum

Date:	03 November 2016		
То:	Bob Anderson		

Cindy Bartlett

From: Mary Tyler

CC: J. Caprio

Subject: Stage 2A Data Validation - Level II Data Deliverable – Eurofins Eaton Analytical Report Number 617739

SITE: PNG0714

INTRODUCTION

This report summarizes the findings of the Stage 2A data validation of seven water samples, one field duplicate sample and one trip blank, collected October 17, 2016, as part of the project PNG0714 sampling event. The analyses were performed at Eurofins Eaton Analytical, Monrovia, California. The samples were prepared and analyzed by the following analytical tests:

- Perfluorinated alkyl acids (PFAAs) by EPA Method 537
- Selected Metals (Calcium, Magnesium, Potassium and Sodium) by EPA Method 200.7
- Anions (Nitrate as N, Nitrate as NO₃, Nitrite as N and total Nitrate, Nitrate-N by Calculation) by EPA Method 300.0

EXECUTIVE SUMMARY

The samples were handled, prepared, and measured in the same manner under similar prescribed conditions.

Based on this Stage 2A data validation covering the quality control (QC) parameters listed below, the data as qualified are usable for meeting project objectives. Qualified data should be used within the limitations of the qualification.

The data were reviewed based on the information and/or guidance provided in the US EPA Contract Laboratory Program National Functional Guidelines for Superfund Organic Methods Data Review, August 2014 (USEPA-540-R-014-002), the US EPA Contract Laboratory Program National Functional Guidelines for Inorganic Superfund Data Review, August 2014 (USEPA-540-
R-013-001), as well as by the pertinent methods referenced by the data package and technical and professional judgment.

The following samples were analyzed in the data set:

Laboratory ID	Sample ID
201610190441	COI-MW01-20161017
201610190442	COI-MW02-2016017
201610190443	COI-MW03-2016017
201610190444	COI-MW03-20161017-DUP
201610190445	COI-MW04-20161017

Laboratory ID	Sample ID
201610190446	COI-MW05-20161017
201610190447	COI-MW06-20161017
201610190448	COI-MW07-20161017
201610190449	TRIP BLANK-20161017

The samples were received at the laboratory within the criteria of 0-6°C. No sample preservation issues were noted by the laboratory.

The following issue was noted with the chain of custody (COC); this did not result in qualification of the data.

• No collection time was listed on the COC for the trip blank. The laboratory assigned a collection time of 0800.

1.0 PERFLUORINATED ALKYL ACIDS

The samples were analyzed for PFAAs per EPA Method 537. The following compounds were reported:

- Perfluorobutanesulfonic acid, PFBS
- Perfluorodecanoic acid, PFDA
- Perfluorododecanoic acid, PFDoA
- Perfluoroheptanoic acid, PFHpA
- Perfluorohexanesulfonic acid, PFHxS
- Perfluorohexanoic acid, PFHxA
- Perfluorononanoic acid, PFNA
- Perfluorooctanesulfonic acid, PFOS
- Perfluorooctanoic acid, PFOA
- Perfluorotetradecanoic acid, PFTA
- Perfluorotridecanoic acid, PFTrDA
- Perfluoroundecanoic acid, PFUnA.

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The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- ✓ Surrogates
- ✓ Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverable Review

1.1 Overall Assessment

The PFAA data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

1.2 <u>Holding Time</u>

The holding times for the PFAA analysis of a water sample are 14 days from collection to extraction and 28 days from extraction to analysis. The holding times were met for the sample analyses.

1.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Two method blanks were reported (batches 944126 and 944831). The PFAAs were not detected in the method blanks above the method-specified one third (1/3) of the method reporting limits (MRLs).

1.4 <u>Matrix Spike/Matrix Spike Duplicate (MS/MSD)</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Batch MS/MSD pairs were reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

1.5 Laboratory Control Sample (LCS)

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Two LCS/LCS duplicate (LCSD) pairs were reported. The recovery and relative percent difference (RPD) results were within the method specified acceptance criteria.

1.6 <u>Surrogates</u>

Acceptable surrogate recoveries were reported for the sample analyses.

1.7 <u>Trip Blank</u>

A trip blank, TRIP BLANK-20161017, was collected with the sample set. The PFAAs were not detected in the trip blank above the MRLs.

1.8 <u>Field Duplicate</u>

A field duplicate sample, COI-MW03-20161017-DUP, was collected with the sample set. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicate and the original sample, COI-MW03-20161017.

1.9 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

1.10 Electronic Data Deliverable (EDD) Review

Results and sample IDs in the EDD were reviewed against the information provided by the associated level II report at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II report and the EDD.

2.0 SELECTED METALS

The samples were analyzed for selected metals per EPA Method 200.7 (Calcium, magnesium, potassium and sodium).

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ✓ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- \checkmark Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverable Review

2.1 Overall Assessment

The metals data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

2.2 <u>Holding Time</u>

The holding time for the metals analysis of a water sample is 180 days from collection to analysis. The holding times were met for the sample analyses.

2.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). One method blank was reported (batch 943898); the method blank was analyzed twice, once in each analytical batch (batches 944503 and 945221). The metals were not detected in the method blank above the MRLs.

2.4 <u>Matrix Spike/Matrix Spike Duplicate</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Four sample set specific MS/MSD pairs were reported, using samples COI-MW01-20161017, COI-MW02-2016017, COI-MW05-20161017 and COI-MW06-20161017. The recovery and RPD results were within the laboratory specified acceptance criteria

2.5 <u>Laboratory Control Sample</u>

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). One LCS/LCSD pair was reported; the LCS/LCSD pair analyzed twice, once in each analytical batch. The recovery and RPD results were within the method specified acceptance criteria.

2.6 <u>Trip Blank</u>

The trip blank was not analyzed for metals.

2.7 <u>Field Duplicate</u>

A field duplicate sample, COI-MW03-20161017-DUP, was collected with the sample set. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicate and the original sample, COI-MW03-20161017.

2.8 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

2.9 <u>Electronic Data Deliverable Review</u>

Results and sample IDs in the EDD were reviewed against the information provided by the associated level II report at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II report and the EDD.

3.0 ANIONS

The samples were analyzed for anions by EPA method 300.0 (Nitrate as N, nitrate as NO₃, nitrite as N and total nitrate, nitrate-N by calculation).

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Final Review: JK Caprio 11/10/16

The areas of data review are listed below. A leading check mark (\checkmark) indicates an area of review in which the data were acceptable. A preceding crossed circle (\otimes) signifies areas where issues were raised during the course of the validation review and should be considered to determine the impact on data quality and usability.

- ✓ Overall Assessment
- ⊗ Holding Time
- ✓ Method Blank
- ✓ Matrix Spike/Matrix Spike Duplicate
- ✓ Laboratory Control Sample
- \checkmark Trip Blank
- ✓ Field Duplicate
- ✓ Sensitivity
- ✓ Electronic Data Deliverable Review

3.1 <u>Overall Assessment</u>

The anion data reported in this package are considered to be usable for meeting project objectives. The results are considered to be valid; the analytical completeness defined as the ratio of the number of valid analytical results (valid analytical results include values qualified as estimated) to the total number of analytical results requested on samples submitted for analysis, for the project is 100%.

3.2 <u>Holding Time</u>

The holding times for water samples are listed below.

Analysis	Holding Time
Nitrate as N, Nitrate as NO ₃ , Nitrite as N and Total Nitrate, Nitrate-N	48 hours from collection to analysis
by calculation by EPA Method 300.0	
Chloride and Sulfate by EPA Method 300.0	28 days from collection to analysis

The holding times were met for the sample analyses, with the following exceptions. The nitrate and nitrite analyses of sample COI-MW02-2016017 were performed 5 minutes outside the 48 hour holding time. Since this sample was analyzed within the hour after 48 hours from collection, no qualifications were applied to the data, based on professional and technical judgment. The nitrate and nitrite analyses of sample COI-MW01-20161017 were performed 5 hours outside the 48 hour holding time. Therefore, based on professional and technical judgment, the nondetect results of nitrate as N, nitrate as NO₃, nitrite as N and total nitrate, nitrate-N by calculation in sample COI-MW01-20161017 were UJ qualified as estimated less than the MRLs.

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Sample ID	Analyte	Laboratory result	Laboratory Flag	Validation result (mg/L)	Validation Qualifier*	Reason Code**
COI-MW01- 20161017	Nitrate (as N)	0.10	UH3	0.10	UJ	2
COI-MW01- 20161017	Nitrate (as NO3)	0.44	UH3	0.44	UJ	2
COI-MW01- 20161017	Nitrite (as N)	0.050	UH3	0.050	UJ	2
COI-MW01- 20161017	Nitrate and Nitrite (as N)	0.10	UH3	0.10	UJ	2

mg/L-milligram per liter

U-not detected at the stated MRL

H3-laboratoary flag indicating that the sample was received and/or analysis requested past the holding time

* Validation qualifiers are defined in Attachment 1 at the end of this report

**Reason codes are defined in Attachment 2 at the end of this report

3.3 <u>Method Blank</u>

Method blanks were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Four method blanks were reported (batches 943714, 943715, 943786 and 943891). The anions were not detected in the method blanks above the MRLs.

3.4 <u>Matrix Spike/Matrix Spike Duplicate</u>

MS/MSD pairs were analyzed at the proper frequency for the number and types of samples analyzed (one pair per batch of 20 samples). Batch MS/MSD pairs were reported. Since these are batch QC, the results do not affect the samples in this data set and qualifications were not applied to the data.

3.5 <u>Laboratory Control Sample</u>

LCSs were analyzed at the proper frequency for the number and types of samples analyzed (one per batch of 20 samples). Four LCS/LCSD pairs were reported. The recovery and RPD results were within the method specified acceptance criteria.

3.6 <u>Trip Blank</u>

The trip blank was not analyzed for anions.

3.7 <u>Field Duplicate</u>

A field duplicate sample, COI-MW03-20161017-DUP, was collected with the sample set. Acceptable precision (RPD \leq 30% for concentrations greater than two times the MRLs, RPD \leq 50% for concentrations within two times the MRLs) was demonstrated between the field duplicate and the original sample, COI-MW03-20161017.

3.8 <u>Sensitivity</u>

The samples were reported to the MRLs. No elevated nondetect results were reported for the samples.

3.9 <u>Electronic Data Deliverable Review</u>

Results and sample IDs in the EDD were reviewed against the information provided by the associated level II report at a minimum of 20% as part of the data validation process. No discrepancies were identified between the level II report and the EDD.

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ATTACHMENT 1 DATA VALIDATION QUALIFIER DEFINITIONS AND INTERPRETATION KEY Assigned by Geosyntec's Data Validation Team

DATA QUALIFIER DEFINITIONS

- U The analyte was analyzed for, but was not detected above the reported sample quantitation limit.
- J The analyte was positively identified; the associated numerical value is the approximate concentration of the analyte in the sample.
- J+ The analyte was positively identified; however, the associated numerical value is likely to be higher than the concentration of the analyte in the sample due to positive bias of associated QC or calibration data or attributable to matrix interference.
- J- The analyte was positively identified; however, the associated numerical value is likely to be lower than the concentration of the analyte in the sample due to negative bias of associated QC or calibration data or attributable to matrix interference.
- UJ The analyte was not detected above the reported sample quantitation limit. However, the reported quantitation limit is approximate and may or may not represent the actual limit of quantitation necessary to accurately and precisely measure the analyte in the sample.
- R The sample results are rejected due to serious deficiencies in the ability to analyze the sample and meet quality control criteria. The presence or absence of the analyte cannot be verified.

ATTACHMENT 2 DATA VALIDATION REASON CODES Assigned by Geosyntec's Data Validation Team

Valid Value	Description
1	Preservation requirement not met
2	Analysis holding time exceeded
3	Blank contamination (i.e., method, trip, equipment, etc.)
4	Matrix spike/matrix spike duplicate recovery or RPD outside limits
5	LCS recovery outside limits
6	Surrogate recovery outside limits
7	Field Duplicate RPD exceeded
8	Serial dilution percent difference exceeded
9	Calibration criteria not met
10	Linear range exceeded
11	Internal standard criteria not met
12	Lab duplicates RPD exceeded
13	Other

RPD-relative percent difference

APPENDIX D 50BS WELL VIDEO SURVEY PHOTO-LOG

GEOSYNTEC CONSULTANTS Photographic Record Geosyntec ^D consultants					
Client:	Oregon DEQ	Project Number: PNG0714			
Subject S	Subject Site: Issaquah Phase II Hydrologic Investigation – Well 5 obs				
Photograp	ph 1		100		
Date:			×.		
N/A			The state of the		
Direction	:	and a state of the second	A SUBARTANIA		
N/A		A1' DA"	and the second		
Comment	ts:	41 04	100 200		
1 st joint at	t 41'04".				
No visible or weldin	g nor intrusion of filter				
pack sand	l. Small amounts of				
iron bacte	eria present.				
D					
Photograp	ph 2		AND MEN TO A		
Date.					
N/A			and the second		
Direction	:				
N/A		and the second			
Comment	ts:	1011 05"	and the second		
4 th Joint a	ut 101'05".		A LARCAR		
or weldin	g nor intrusion of filter				
pack sand	l. Small amounts of				
iron bacte	eria present.				
			the stand		

Is saquah W5 obs Photo Log. docx

	GEOSYNTEC CONSULTANTS Photographic Record	
Client: Oregon DEQ	Project Number: PNG0714	
Subject Site: Issaquah Phase II Hy	drologic Investigation – Well 5 obs	
Photograph 3		
Date:		1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-1-
N/A		
Direction:	Stand Stand Star 19	
N/A	101100	
Comments: 7 th Joint at 181'09"	101 00	
No visible deterioration of joint		
or welding nor intrusion of filter		
iron bacteria present.		
		ALC: ANTERIA
Photograph 4		
Date:		A REAL PROPERTY.
N/A		a the same
Direction:		Sec. 1
N/A		
Comments:	281'11"	
12 th Joint at 281'11". No visible deterioration of joint		e terter
or welding nor intrusion of filter		
pack sand. Small amounts of iron bacteria present.		harden im 200
non oueterne present.		

Is saquah W5 obs Photo Log. docx

GEOSYNTEC CONSULTANTS Photographic Record Geosyntec ^D			
Client: Oregon DEQ	Project Number: PNG0714		
Subject Site: Issaquah Phase II Hy	drologic Investigation – Well 5 obs		
Photograph 5	and the second se		
Date:			
N/A			
Direction:			
N/A	304' 10"		
Comments:		1. C.	
Concrete plug at 304'10".			
Location above well screen. No	A Contract		
seepage through/around			
concrete.			
	the second s	STATES -	