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Release Fraction Evaluation

J. A. Bamberger J. A. Glissmeyer

January 2004

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Pacific Northwest National Laboratory Richland, Washington 99352

Summary

This document presents results of experiments conducted to measure release fractions during certain tank retrieval processes. The tests were performed in a $\frac{1}{4}$ - scale model of a 75-ft (22.9-m) diameter million gallon (3785 m³) waste storage tank. The retrieval processes simulated were:

- 1. Discharging liquid or slurry from the mouth of a vertically oriented 2-in. Schedule 40 pipe [ID-2.067 in. (5.25 cm)]. The discharging material was in free-fall from the mouth of the pipe near the top of the tank into a liquid or slurry pool at the bottom of the tank, a distance of 11.4 ft (3.48 m).
- 2. The jet from a 9/16-in. (1.43 cm-) diameter nozzle transferring liquid or slurry waste from one side of the tank to the other. The discharging liquid was aimed at the opposite side of the tank from the nozzle and either impacted the tank wall or fell into a liquid or slurry pool in the bottom of the tank.
- 3. A high pressure fan jet of liquid striking a steel plate or simulated waste from a stand-off distance of a few inches.

For each process, a water-soluble fluorescent dye was added to the liquid fraction as a tracer. Kaolin clay was used to represent the solids. The tank was covered and there was no forced ventilation in the tank during the tests.

Six air samples were collected during each test. The air samples were collected at fixed positions in the tank. The air sample filters were dried and weighed to determine the solids collection. The fluorescent dye was then leached from each filter and quantified with a fluorometer to determine the collection of liquid. Samples of the slurry and liquid simulants were also collected to determine the quantities of simulant used in each test.

To calculate the release fraction, the quantity collected on each air sample was adjusted for the fraction of the tank volume sampled and divided by the quantity of material exposed in the simulation. The method was not as sensitive for the solids content as it was for the liquid content; but in those instances where a solids release fraction was determined, it was in agreement with that of the liquid phase.

Release fractions are commonly used to make conservative estimates of emissions from processes. Usually, rather gross assumptions are made in such estimates, such as the total failure of abatement equipment and the use of maximum inventory values. Consequently, it is common practice to report bounding release fraction values with single digit accuracy. The release fractions for the top of the unventilated tank ranged from 9 x 10^{-7} to 8 x 10^{-5} depending on the process simulated. The particle size distribution was determined to be log-normally distributed with geometric mean diameters ranging from 3 to 8 µm. Thus, all particles are conservatively considered "respirable."

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1.0 Introduction

Waste mobilization and retrieval processes are implemented to remediate nuclear waste stored in 75-ft (22.9-m) diameter million-gallon (3785-m³) underground storage tanks at the Hanford Site in Washington state. These processes are developed based on the characteristics of the waste to be retrieved and other operational constraints. These processes have the potential to generate aerosols or suspend solids during operation. The purpose of this study was to evaluate the aerosol generation performance of several types of fluid transport processes that could be considered for future implementation. These processes included:

- 1. Discharging liquid or slurry from the mouth of a vertically oriented two-inch pipe. The discharging material was in free-fall from the mouth of the pipe near the top of the tank into a liquid or slurry pool at the bottom of the tank, a distance of 11.4 ft (3.48 m).
- 2. The jet from a nozzle transferring liquid or slurry waste from one side of the tank to the other. The discharging liquid was aimed at the opposite side of the tank from the nozzle and either impacted the tank wall or fell into a liquid or slurry pool in the bottom of the tank.
- 3. A high pressure fan jet of liquid striking a steel plate or simulated waste from a stand-off distance of a few inches.

To characterize the aerosol a water-soluble fluorescent dye was added to the liquid fraction as a tracer. Kaolin clay, added to the liquid fraction, was used to represent the solids. The tank was covered, and there was no forced ventilation in the tank during the tests. To calculate the release fraction, the quantity collected on each air sample was adjusted for the fraction of the tank volume sampled and divided by the quantity of material exposed in the simulation. These experiments build on research conducted at PNNL in the 1980s to characterize aerosols generated by releases of powders and solutions (Ballinger et al 1986, 1987, 1988 and Sutter et al 1981).

2.0 Conclusions and Recommendations

2.1 Conclusions

The release fraction results for this study are listed in Table 2.1. This summary focuses on the values obtained from measurements taken at the top of the tank which is the location from which ventilation air may be drawn. The liquid fraction results are presented because this method of measurement (dye tracer) was more sensitive than that used for the solids fraction (solids loading). These results range over nearly two orders of magnitude. It is noted that the height of the test tank is lower than that of the full scale tank by a factor of 4, but the tests were not conducted to provide the release fraction as a function of height.^{a)} The release fractions calculated based on samples obtained near the point of the process or fluid stream ranged from the same to as much as an order of magnitude higher than the release fractions obtained from measurements taken near the tank top at an elevation of 11 ft (3.35 m).

Run	Process	Release Fraction at Scaled Tank Top
	Pipe Transfer	
2	Liquid discharge at low flow rate	0.9 x 10 ⁻⁶
11	Slurry discharge at low flow rate	2 x 10 ⁻⁶
12	Slurry discharge at high flow rate	4 x 10 ⁻⁶
	Transfer Jet	
5	Liquid discharge	3 x 10 ⁻⁶
10	Slurry discharge	2 x 10 ⁻⁶
	Cutting Jet	
8	Dislodging simulant at low pressure	8 x 10 ⁻⁵
9	Dislodging simulant at high pressure	8 x 10 ⁻⁵

2.2 Recommendations

The release fraction measurements could be reported with all significant digits; however, there are simplifications made in simulating processes and significant spatial and orientation differences between the individual sampling locations. Release fractions are commonly used to make conservative estimates of emissions from processes. Usually, rather gross assumptions are made in such estimates, such as the

^{a)} Height influences release fractions in two competing ways: 1) greater height of a spill is likely to increase aerosol generated, and 2) greater height from the source of aerosol of generation to the potential receptor decreases concentration due to particle depletion mechanisms.

total failure of abatement equipment and the use of maximum inventory values. Consequently, it is common practice to report bounding release fraction values with single digit accuracy.

3.0 Aerosol Generation Processes

Three processes were identified as potential sources of aerosol generation during their in-tank operation. These processes were investigated to determine their release fractions.

3.1 Cutting Jet

Cutting jets are mounted on an articulated arm or crawler and are used for cutting, dislodging, or cleaning solids or slurry from surfaces. The jet configuration to be used in waste retrieval equipment includes five jets mounted in a semicircle directed downward. The jets are commercially available wash jets (Spraying Systems Company^{a)} ¹/₄ MEG 2505). Tests conducted at the Hanford cold test facility by others showed that the preferred pressure for cutting jet operation is 300 psig (2.07 MPag). The proposed jet system can operate at pressures up to 1200 psig (8.27 MPag). The effective stand-off distance for the cutting jets ranges from 1 to 5 in. based on the deployment configuration. The fan jet has a spray angle of 25 degrees. For this evaluation a single jet mounted in line with a pressure gage was used as shown in Figure 3.1.



Figure 3.1. Cutting jet nozzle configuration

3.2 Transfer Jet

The transfer jet nozzle is mounted on the crawler as shown in Figure 3.2. The transfer nozzle configuration tested was mounted on the side of the tank and is shown in Figure 3.3. The nozzle operates at relatively low flow rates and pressures and is used to spray fluid from the crawler location to a collection location near a retrieval pump or other retrieval device.

^{a)} Spraying Systems Company, Wheaton, Illinois.



Figure 3.2. Crawler showing transfer nozzle



Figure 3.3. Transfer jet nozzle configuration

3.3 Tank Transfer: Flow from an Open Pipe

Transfers of fluid or slurry from tank to tank occur through nominal 2 in. (5 cm) diameter pipes or 2 in. (5 cm) hose in hoses. Some transfers may be conducted using nominal 3 in. (7.6 cm) diameter pipe. The flow rate range is 50 to 60 gpm (.0032 to 0.0038 m^3 /s) for hose in hose transfers, 50 to 100 gpm (0.0032 to 0.0063 m^3 /s) for 2 in. pipe transfers, and 50 to 160 gpm (0.0032 to 0.0101 m^3 /s) for 3 in. (7.6 cm) pipe transfers. The nozzle configuration evaluated is shown in Figure 3.3.



Figure 3.4. Pipe transfer nozzle configuration used to model flow from tank transfers

3.4 Aerosol Generation Experiment Configuration

All aerosol generation characterization tests were conducted in the ¼-scale tank located in the 336 bldg. of Hanford's 300 Area to provide an enclosed structure (Bamberger et al 1993). The ¼-scale tank is a ¼-scale model of a 75-ft (22.9 m) 1 million gallon (3785 m³) Hanford double-shell tank. Three segments of the ¼-scale tank dome were installed; the remaining quadrant of the tank was closed with a plastic tarp. The enclosure prevented aerosol from entering or leaving the tank test chamber. Fluid or slurry was pumped from the mix tank through the nozzles into the ¼-scale tank. Figure 3.3 shows the arrangement of the ¼-scale tank with the top rim of the tank, part of the dome and one of the test nozzles. The air samplers clustered near the top of the tank are shown in Figure 3.4. The locations of the experiment components are summarized in Table 3.1.

The cutting jet was powered using a portable gasoline driven pressure washer. The unit provides pressure up to 3200 psig (22.1 MPag). The unit was instrumented with pressure gages to measure the pressure at the unit and at the nozzle. The pressure washer was located outside of the building away from the roll-up door so that no gasoline exhaust fumes from operation could blow into the building.

An Ingersol Rand Standard Pump Division centrifugal pump was used to circulate fluid between the mix tank and the nozzles being evaluated in the test tank. The pump has a capacity of 150 gpm ($0.0095 \text{ m}^3/\text{s}$) and 275 psig (1.90 MPag). The mix tank supplies fluid to this pump. After the mix tank is emptied, the fluid or slurry is pumped back into the mix tank to provide the desired number of operating cycles. A

compressed air driven transfer pump was used to move fluid from the $\frac{1}{4}$ -scale tank to the mix tank. The transfer pump is powered by a rented 1650 scfm (0.779 sm³/s), diesel driven, portable air compressor located outside on the east side of the building.

A digital video camera mounted on the tank walkway around the perimeter of the tank was used to film the jet during the tests.

		Coordinate							
Item	Location	Radial		Angular	Eleva	tion			
		R=0 at ta	nk center	ુ =0 at North	Z=0 a	t tank floor			
Sampler		ft	m	degrees	ft	m			
Sampler 1	Top of tank	1	0.31	188	11.5	3.51			
Sampler 2	Top of tank	1	0.31	230	11	3.35			
Sampler 3	Top of tank	1	0.31	230	11	3.35			
Sampler 4	Side of tank	9.4	2.87	158	5	1.52			
Sampler 5	Side of tank	9.4	2.87	186	5	1.52			
Sampler 6	In tank space	4.6	1.40	155	4	1.22			
Dust Monitor	Top of tank	1	0.31	188	11.5	3.51			
Optical Particle	Top of tank	1	0.31	188	11	3.35			
Counter (OPC)									
Process Hardware	Orientation								
Discharge pipe	Vertical	8.5	2.60	8	11.4	3.48			
Transfer jet	Horizontal	8	2.44	175	8	2.44			
Cutting jet	Vertical	4	1.22	153	1	3.35			
Tank Dimensions									
Tank height					8	2.44			
Dome height					3.5	1.07			
Riser height					0.5	0.15			
Total height					12	3.66			
Tank radius		9.38	2.86						
Domed Area				188.5 to 360 to					
				98.5					
Plastic Covered				98.5 to 188.5					
Area									

 Table 3.1.
 Locations of hardware and sampling equipment

4.0 Measurement Methods

Variables monitored during the tests included:

- Fluid initial and final level in the mix tank (used to calculate the volume of fluid used per cycle)
- Pump operating time used for calculating the average flow rate per run
- Real-time aerosol concentration
- Real-time aerosol concentration as a function of particle size
- Liquid and solid aerosol concentration measured with integrating air samplers.

Twelve test runs were conducted. The first run provided background measurements. The rest covered the range of operating parameters for the three simulated processes.

4.1 Aerosol Measurement Methods

Three methods were used for monitoring air concentration in the ¹/₄-scale tank during the test runs. The first was to collect air samples on filters. The second method was to qualitatively monitor particle size and concentration in near real time with an optical particle counter (OPC).^{a)} A third method, using a real-time dust monitor, was unsuccessful and produced no meaningful data. The cluster of air monitors located at the top of the tank is shown in Figure 3.4, and Figure 4.1 shows the location of Air Sampler 6 inside the tank. Appendix A contains the detailed sampling and analysis procedure. The methods are further described below.



Figure 4.1. Location of air sampler 6 inside tank

^{a)} Model 2408A, Pacific Scientific Instruments (formerly Met-One), Grants Pass, Oregon.

4.1.1 Air Sampling

Samples of the airborne material were collected during the background run and eight of the process simulations. The samples were collected on 25–mm diameter mixed cellulose ester membrane filters.^{a)} The samples were contained in open-face filter holders^{b)} which were connected by flexible tubes to personal sampling pumps^c located outside the tank.

The filter holders were placed to collect samples at four locations in the tank. Three (samplers 1, 2, and 3) were collocated at the center of the top of the tank. Two (samplers 4 and 5) were located along the tank wall 9.4 ft (2.87 m) from the tank center at elevations of 8 (2.44 m) and 5 ft (1.52 m) above the tank floor. The sixth air sampler was in the central area of the tank and in close proximity to the process stream and at an elevation of 5 ft (1.52 m) from the tank floor. The sampler positions were listed in Table 3.1. Only the three air samplers at the top of the tank were expected to measure aerosol at a location approximating that where exhaust air is drawn from the full scale tanks.

At the beginning of each test day, another filter holder was prepared and used to collect a background air sample. It was operated at sampling position 4. Sampling flow rates were 0.75 lpm (liters per minute) for the sampler near the process stream and 1.5 lpm for the others. A calibrator^d was used to set the flow rates at the sampling pumps prior to and after the series of measurements.

The filter holder consisted of a filter cassette and the outer holder. Before a test run, these parts were thoroughly cleaned with water and sonicated in deionized water. The components were dried, and the cassettes were loaded with a filter disk. The cassettes were tare weighed, and the assembled filter holders were packaged for transport. Two cassettes were included with each run as field blanks.

Most test runs included several sequential operations of the simulated process. To provide the desired operating time, fluid in the mix tank was sprayed into the test tank. When the mix tank was empty, the spray process was stopped, and the fluid was pumped back into the mix tank. This process was completed up to four times to provide the desired operating time. The air samplers were always operated for the entire duration of the test run. At the end of a run, the air samplers were retrieved and packaged for transport. In the lab, the cassettes were removed from the filter holders and dried for thirty minutes in an oven at about 90°F (32 °C). The cassettes were allowed to equilibrate to room temperature and were reweighed. For analysis, the filters were removed from the cassettes and placed in known quantities of deionized water to determine the fluorescent dye content. After several hours, the water was analyzed for the content of fluorescent dye as compared to prepared standards. Each run was analyzed as a batch. Standards, field blanks, and process blanks were analyzed with each batch of air samples.

Air sampler tare weight, final weight, flow rate, sampling duration, and analytical data were recorded on data sheets. The hand written data were then transferred to spreadsheets.

^{a)} MF-Millipore, Model HAWP 02500, Millipore Inc., Billerica, Massachusetts.

^{b)} SKC Incorporated, IOM Inhalable Particle Sampler, Eighty Four, Pennsylvania. This sampler was developed at the Institute of Occupational Medicine, Edinburgh, UK by Mark and Vincent (1986)

^{c)} SKC Incorporated, Model 224, Eighty Four, Pennsylvania.

^{d)} DC-Lite 717-04, SKC Incorporated, Eighty Four, PA.

4.1.2 Analysis of Fluorometry Results

Air samplers were operated during eight test runs and one background run. The estimated aerosol release fraction was calculated for each air sample according to Equation 4.1. The dye in the sample is obtained from the fluorometry results for each sample. The dye in the process is obtained from analyses of process water samples and measurements of the process water as listed in Section 5. The tank volume was calculated from tank geometry, and the sample volume was calculated from the fluorometry results and release fraction calculations are tabulated in Appendix C.

Release fraction of liquid =
$$\frac{\text{dye in sample}}{\text{dye in process}} \bullet \frac{\text{tank volume}}{\text{air sample volume}}$$
 Eq. 4.1

4.1.3 Analysis of Dry Weight Results

The balance used was accurate to tenths of a milligram. When all of the background air samples and field blanks were considered, weight changes ranged from 0.1 to -0.5 mg; and 53% of these samples showed 0.0 mg weight change during a run. The mean weight change of these samples was -0.09 mg. Only a sample weight gain of 0.2 mg (200µg) was considered as a valid measurement of deposited solids for this report.

After drying the samples, any sample weight gain should be attributed to the solids fraction of the waste simulant and the soluble fluorescent dye. However, the content of fluorescent dye measured by fluorometry on sample filters ranged from 7.3 μ g (Run 7, sample 7B4) to 3.2 ng (Run 1, sample 1A4). Consequently, the contribution of the fluorescent dye was ignored in the weight gain measurements.

Solids were introduced into the process simulations during Runs 8 through 12. For these runs, the air sample weight gain should represent the release of the solids portion of the process slurry. Solids were selected based on nonhazardous simulants developed to model performance of equipment planned for use in radioactive waste tanks (Powell, Golcar, Geeting 1997). The simulant used was selected for evaluation of the process equipment described in Section 3.

The equation to calculate the solids release fractions is given by Equation 4.2. The calculations of dry solids release fractions are located in Appendix C.

Release fraction of solids = $\frac{\text{solids in sample}}{\text{solids in process}} \bullet \frac{\text{tank volume}}{\text{air sample volume}}$ Eq. 4.2

4.2 **Optical Particle Counter**

The OPC categorizes particulate into one of six size brackets based on the amount of laser light scattered by a particle that is equivalent to that of a spherical calibration particle made of polystyrene latex. To

complicate analysis, the OPC has limited range, and there are inlet losses that are a function of particle size. However, there is value in using the particle size data from the OPC to observe trends.

Each OPC data record is the total particle count in each size range for a one-minute interval. There was a one second lag time between samples. The OPC aerosol inlet used was the manufacturer's inverted cone nozzle which was attached to the instrument with flexible conductive tubing to inhibit particle plate out in the tube. The inlet was collocated with air samplers 2 and 3 at the center of the top of the tank, and the OPC was 2 ft (0.61 m) above that and outside the enclosed air space. The sampling flow rate was 28.3 lpm. There was a single bend in the flexible tubing. The OPC was operated throughout each testing day and for all test runs.

5.0 Experiments

The experimental test matrix evaluated three processes: cutting jet, transfer jet, and flow from an open pipe, and two fluids: water and slurry. In addition, the cutting jet impacted two surfaces: steel and simulant. The processes were operated at several flow rates.

5.1 Water and Slurry Tests

The water (Runs 2 through 7) and slurry (Runs 10 through 12) spray tests were conducted based on the following procedure. Run 1 was a background run. Fluorescent dye was added to the water in the mix tank to be used as a tracer for identifying the amount of water captured on the air samplers. To obtain slurry, water was sprayed through the cutting jet into a container of simulant during Runs 8 and 9. The slurry was used as the starting fluid during Runs 10 through 12. The solid captured on filters was measured to determine the solids in suspension. The procedure during the water and slurry tests included:

- Configure piping to include the process hardware that is to be tested.
- Set the valves to provide the desired flow rate.
- Record the liquid level in the mix tank.
- Turn on aerosol sampling equipment.
- Turn on video camera.
- Start the pump to route fluid from the mix tank through the process hardware.
- Observe the spray produced.
- Record the time the pump stopped and final level in mix tank.

Note: Based on the capacity of the mix tank, experiments to evaluate the transfer jet and the tank transfer were conducted as a series of sequential runs to provide an adequate sampling time. For these evaluations, the following additional steps were followed.

- Pump the liquid from the ¹/₄-scale tank to the mix tank.
- Record the level in the mix tank and complete the steps listed above as many times as required to provide the desired operating time.
- Stop sampling pumps.
- Remove samplers and analyze the samples.

5.2 Cutting Jet Tests

The following procedure was used for the cutting jet tests for impacting steel or for erosion of simulant.

- Place the desired surface beneath the jet nozzle: either a steel plate or a container of simulant. The container will have 1 in. vertical slits on either side to permit water to run out of the container.
- Position the container so it can be moved beneath the wash jet.
- Configure piping for the cutting jet.
- Set the valves to provide the approximate flow rate.
- Record the liquid level in the mix tank.
- Turn on aerosol sampling equipment.
- Turn on video camera.

- Start the pressure sprayer.
- If simulant is being used, incrementally move the simulant container beneath the wash jet during the test period.
- Observe the spray produced.
- Record time pump stopped and final level in mix tank.
- Use a new container of simulant for each cutting jet test.

5.3 Cutting Jet Aerosol Generation Tests

The tests with the cutting jet are described first, because the cutting jet was used to produce the slurry used during solids suspension tests conducted with the other processes. Each test run is described in the following subsections.

5.3.1 Run 7: Cutting Jet Impacting Steel Plate at Low Pressure

These tests were conducted by pumping water with fluorescent dye through the cutting jet which impacted a steel plate, mounted above the water level in the tank. Each test consisted of a single operating cycle. The fluid volume and flow rate data are summarized in Table 5.1. Photographs taken during the sequence are shown in Figure 5.1. The jet operating pressure was in the 340 to 380 psig (2.34 to 2.62 MPag) range. The monitored test conditions are fluid volume and time which provide data to calculate the average flow rate. The cutting jet may be used to scarify solid waste that is otherwise not sufficiently mobilized by other dislodging mechanisms. If the water-based cutting jet actually strikes bare steel, it is not striking waste, and waste is not being released at that particular time. Consequently, a measured release fraction from this run is not actually applicable to waste retrieval.



Table 5.1. Run 7 flow rate data through the cutting jet

Cycle 7-1 Figure 5.1. Photos from Run 7 showing the jet pattern on the steel plate

5.3.2 Run 8: Cutting Jet Impacting Simulant at Low Pressure

These tests were conducted by pumping water with fluorescent dye through the cutting jet which impacted simulant contained in a box mounted above the water level in the tank. Each test consisted of one cycle. The fluid volume and flow rate data are summarized in Table 5.2. Photographs taken during the sequence are shown in Figure 5.2. Every 5 minutes during the 30 minute test, the simulant position was incremented to move new simulant beneath the jet. The jet operating pressure was in the 300 psig (2.07 MPag) range. During the run the jet dislodged 9 kg of simulant out of 36.3 kg in the container. The monitored test conditions are fluid volume and time which provide data to calculate the average flow rate.

Run	Cycle	Time	Change in 1	Height	ht Fluid Volume Average		Average	Flow Rate	
8	1	sec 1806	in. 7 75	cm 19.7	gal 39.1	m^{3} 0.1478	gpm 1 30	m³/s	

 Table 5.2
 Run 8 flow rate data through the cutting jet



Cycle 8-1



5.3.3 Run 9: Cutting Jet Impacting Simulant at High Pressure

These tests were conducted by pumping water with fluorescent dye through the cutting jet which impacted simulant contained in a box mounted above the water level in the tank. Each test consisted of one cycle. The fluid volume and flow rate data are summarized in Table 5.3. Photographs taken during the sequence are shown in Figure 5.3. Every 5 min during the 30 min test, the simulant position was incremented to move new simulant beneath the jet. The jet operating pressure was in the 1200 psig (8.27 MPag) range. During the run the jet dislodged 24 kg out of 34.1 kg of simulant in the container. The monitored test conditions are fluid volume and time which provide data to calculate the average flow rate.

Run	Cycle	Time	Change in Height		Fluid	l Volume	Average Flow Rate	
		sec	in.	cm	gal	m ³	gpm	m ³ /s
9	1	1804	16.25	41.3	81.9	0.3100	2.7	0.000172





Cycle 9-1

Figure 5.3. Photos from Run 9 showing the cutting jet impacting the sludge simulant at high pressure

5.4 Tank Transfer Aerosol and Solids Generation Tests

Aerosol generation tests were conducted with water for Runs 2, 3, and 4 and slurry for Runs 11 and 12. The slurry was generated by adding simulant to the water in the mix tank. The mixer was operated to keep the solids in suspension. Descriptions and a series of pictures taken during each of the runs follows.

5.4.1 Run 2: Water at Low Flow Rate

These tests were conducted by pumping water with fluorescent dye through the 2 in. diameter nozzle. Each test consisted of four cycles. The operating time, fluid volume and calculated average flow rates are summarized in Table 5.4. Photographs taken during the sequence are shown in Figure 5.4.

Run	Cycle	Time	Change in Height		Fluid	Volume	Average	Flow Rate
		sec	in.	cm	gal	m ³	gpm	m ³ /s
2	1	374	37.5	95.3	189.0	0.7154	30.3	0.001913
	2	312	37.5	95.3	189.0	0.7154	36.3	0.002293
	3	401	37.5	95.3	189.0	0.7154	28.3	0.001784
	4	257	37.5	95.3	189.0	0.7154	44.1	0.002784
	Mean Std.	336	37.5	95.3	189.0	0.7154	34.8	0.002193
	Dev.	64.5	0	0.0	0.0	0.0	7.1	0.000449
					Total l	Fluid Spra	ayed	
					gal	m ³		
					755.9	2.862		

Table 5.4. Run 2 flow rate data through a discharge pipe at low flow rate



Cycle 2-3



Cycle 2-4

Figure 5.4. Photos from Run 2 showing water transfer from a discharge pipe at low flow rate

5.4.2 Run 3: Water at High Flow Rate

These tests were conducted by pumping water through the 2 in. diameter nozzle. Each test consisted of four cycles. The operating time, fluid volume and calculated average flow rates are summarized in Table 5.5. Photographs taken during the sequence are shown in Figure 5.5.

Run	Cycle	Time	Change in Height		Fluid '	Volume	Average F	low Rate
		sec	in.	cm	gal	m ³	gpm	m ³ /s
3	1	110	38.5	97.8	194.0	0.7345	105.8	0.006677
	2	109	38.5	97.8	194.0	0.7345	106.8	0.006738
	3	111	37.5	95.3	189.0	0.7154	102.2	0.006445
	4	110	37.5	95.3	189.0	0.7154	103.1	0.006504
	Mean Std.	110	38.0	96.5	191.5	0.7249	104.5	0.006591
	Dev.	0.82	0.6	1.5	2.9	0.0	2.2	0.000139
					Total I	Fluid Spra	ayed	
					gal	m ³		
					766.0	2.900		

Table 5.5. Run 3 flow rate data through a discharge pipe at high flow rate

Cycle 3-2

Figure 5.5. Photos from Run 3 showing water transfer from a discharge pipe at high flow rate

5.4.3 Run 4: Water at Highest Flow Rate

These tests were conducted by pumping water through the 2 in. diameter nozzle. Each test consisted of four cycles. The operating time, fluid volume and calculated average flow rates are summarized in Table 5.6. Photographs taken during the sequence are shown in Figure 5.6.

Table 5.6.	Run 4	flow rate	data	through	a discharge	pipe	at highest	flow rate

Run	Cycle	Time	Change in Height		Fluid Volume		Average Flow Rate			
		sec	in.	cm	gal	m ³	gpm	m³/s		
4	1	64	37.5	95.3	189.0	0.7154	177.2	0.011178		
	2	64	37.5	95.3	189.0	0.7154	177.2	0.011178		
	3	61	37.5	95.3	189.0	0.7154	185.9	0.011728		
	4	65	37.5	95.3	189.0	0.7154	174.4	0.011006		
	Mean Std.	63.5	37.5	95.3	189.0	0.7154	178.7	0.011272		
	Dev.	1.73	0.0	0.0	0.0	0.0	5.0	0.000314		
				Total Fluid Sprayed						
					gal	m ³				
					755.9	2.862				

Cycle 4-2

Cycle 4-3

Cycle 4-4 Close up of jet

Figure 5.6. Photos from Run 4 showing water transfer from a discharge pipe at highest flow rate

5.4.4 Run 11: Slurry Transfer at Low Flow Rate

These tests were conducted by pumping slurry through the 2 in. diameter nozzle. Each test consisted of four cycles. The operating time, fluid volume and calculated average flow rates are summarized in Table 5.7. Photographs taken during the sequence are shown in Figure 5.7.

Table 5.7. Run 11 slurry flow rate data through discharge pipe at low flow	rate
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Run	Cycle	Time	Change in Height		Fluid	Volume	Average Flow Rate		
	-	sec	in.	cm	gal	m ³	gpm	m^3/s	
11	1	399	37.5	95.3	189.0	0.7154	28.4	0.001793	
	2	449	37.5	95.3	189.0	0.7154	25.3	0.001593	
	3	211	37.5	95.3	189.0	0.7154	53.7	0.003390	
	4	174	37.5	95.3	189.0	0.7154	65.2	0.004111	
	Mean Std.	308.3	37.5	95.3	189.0	0.7154	43.1	0.002722	
	Dev.	136	0.0	0.0	0.0	0.0	19.4	0.001227	
			Total Fluid Sprayed						
					gal	m ³			
					755.9	2.862			

Close up of jet during cycle 11-4

Figure 5.7. Photos from Run 11 showing slurry transfer through a discharge pipe at low flow rate

5.4.5 Run 12: Slurry Transfer at High Flow Rate

These tests were conducted by pumping slurry through the discharge pipe. Each test consisted of four cycles. The fluid volume and flow rates are summarized in Table 5.8. Photographs taken during the sequence are shown in Figure 5.8.

Run	Cycle	Time	Change in Height		Fluid Volume		Average Flow Rate		
		sec	in.	cm	gal	m ³	gpm	m ³ /s	
12	1	93	37.5	95.3	189.0	0.7154	121.9	0.007692	
	2	92	37.5	95.3	189.0	0.7154	123.3	0.007776	
	3	90	37.5	95.3	189.0	0.7154	126.0	0.007949	
	4	89	37.5	95.3	189.0	0.7154	127.4	0.008038	
	Mean Std.	91	37.5	95.3	189.0	0.7154	124.6	0.007864	
	Dev.	1.8	0.0	0.0	0.0	0.0	2.5	0.000158	
			Total Fluid Sprayed						
					gal	m ³			
					755.9	2.862			

Table 5.8. Run 12 slurry flow rate data through a discharge pipe at high flow rate

Cycle 12-1

Figure 5.8. Photos from Run 12 showing slurry transfer through a discharge pipe at high flow rate

5.5 Transfer Jet Aerosol Generation Tests

5.5.1 Run 5: Water at Low Flow Rate

These tests were conducted by pumping water with fluorescent dye through the 9/16 in. (1.43 cm) diameter transfer jet. Each test consisted of four cycles. The operating time, fluid volume and calculate average flow rates are summarized in Table 5.9. Photographs taken during the sequence are shown in Figure 5.9.

Run	Cycle	Time	Change in Height		Fluid Volume		Average Flow Rate			
		sec	in.	cm	gal	m ³	gpm	m ³ /s		
5	1	414	37.5	95.3	189.0	0.7154	27.4	0.001728		
	2	311	37.5	95.3	189.0	0.7154	36.5	0.002300		
	3	300	37.5	95.3	189.0	0.7154	37.8	0.002385		
	4	417	37.5	95.3	189.0	0.7154	27.2	0.001716		
	Mean	360.5	37.5	95.3	189.0	0.7154	32.2	0.002032		
	Stu. Dev.	63.7	0.0	0.0	0.0	0.0	5.7	0.000360		
				Total Fluid Sprayed						
					gal	m ³				
					755.9	2.862				

 Table 5.9.
 Run 5 flow rate data through a transfer jet at low flow rate

Cycle 5-3

Cycle 5-4

Figure 5.9. Photos from Run 5 showing water transfer jet test sequence at low flow rate

5.5.2 Run 6: Water at High Flow Rate

These tests were conducted by pumping water with fluorescent dye through the 9/16 in. (1.43 cm) diameter transfer jet. Each test consisted of four cycles. The operating time, fluid volume and calculated average flow rates are summarized in Table 5.10. Photographs taken during the sequence are shown in Figure 5.10.

Run	Cycle	Time	Change in Height		Fluid Volume		Average Flow Rate		
		sec	in.	cm	gal	m ³	gpm	m ³ /s	
6	1	201	37.5	95.3	189.0	0.7154	56.4	0.003559	
	2	199	37.5	95.3	189.0	0.7154	57.0	0.003595	
	3	200	37.5	95.3	189.0	0.7154	56.7	0.003577	
	4	199	37.5	95.3	189.0	0.7154	57.0	0.003595	
	Mean Std.	199.8	37.5	95.3	189.0	0.7154	56.8	0.003581	
	Dev.	0.96	0.0	0.0	0.0	0.0	0.3	0.000017	
				Total Fluid Sprayed					
					gal	m ³			
					755.9	2.862			

Table 5.10. Run 6 flow rate data through a transfer jet at high flow rate

Cycle 6-1

Cycle 6-4

Figure 5.10. Photos from Run 6 showing water transfer jet test sequence at high flow rate

5.5.3 Run 10: Slurry at Low Flow Rate

These tests were conducted by pumping slurry with fluorescent dye through the 9/16 in. (1.43 cm) diameter transfer jet. Each test consisted of four cycles. The operating time, fluid volume and calculated average flow rates are summarized in Table 5.11. This test was stopped after the nozzle became plugged during cycle 4. Photographs taken during the sequence are shown in Figure 5.11.

Run	Cycle	Time	Change in Height		Fluid Volume		Average Flow Rate			
		sec	in.	cm	gal	m ³	gpm	m ³ /s		
10	1	525	37.5	95.3	189.0	0.7154	21.6	0.001363		
	2	456	37.5	95.3	189.0	0.7154	24.9	0.001569		
	3	440	37.5	95.3	189.0	0.7154	25.8	0.001626		
	4	537	37.5	95.3	189.0	0.7154	21.1	0.001332		
	Mean	489.5	37.5	95.3	189.0	0.7154	23.3	0.001472		
	Std. Dev.	48.6	0.0	0.0	0.0	0.0	2.3	0.000147		
				Total Fluid Sprayed						
					gal	m ³				
					755.9	2.862				

Table 5.11 .	Run	10 slurr	y flow	rate data	through	a transfer	jet at l	low flow rate
					0		5	

Cycle 10-1

Cycle 10-2

Cycle 10-3

Cycle 10-4

Plugged nozzle at end of cycle 4

Figure 5.11. Photos from Run 10 showing slurry transfer jet test sequence at low flow rate

6.0 Results

This section describes the characteristics of the aerosol created by the processes simulated in the ¹/₄-scale tank. The results presented and discussed include the real-time aerosol concentration, particle size distribution, and the release fractions.

6.1 Aerosol Concentration Results

With each test run, the optical particle counter (OPC) data showed a sharp increase in aerosol concentration from the background level and a gradual decline in aerosol concentration at the conclusion of the process simulation. The particle count data were converted to the volume (or mass) of aerosol in each particle size interval. Plots of these converted OPC data for all of the runs are included in Appendix B.

Figure 6.1. Example plot of real-time aerosol concentration data

Figure 6.1 shows the OPC particle concentration data for Runs 3 and 4 combined. In these runs, the 2-in. nozzle was used to transfer liquid into the tank from the top and the liquid impacted floor or liquid at the bottom of the tank. The operation was repeated eight times, and the airborne concentration peaked each time. The peaks and the baselines in between operations gradually increased. These data show that when a process is repeated rapidly enough the concentration of airborne material continues to ratchet upward, and the concentration does not decrease back to background during the cycles, either because the tank was not ventilated or because sufficient time had not elapsed in between operations.

The total volume concentration peaked at sample interval 80 during the last repetition of the operation. The concentration had not returned to background in this unventilated tank even after 35 sample intervals (about 36 minutes). In Table 6.1, the number of sample intervals (each about 61 seconds) that elapsed from the peak of each runs' final operation to the return to background levels are tabulated. The shortest concentration decay to background observed was 51 samples (about 52 minutes). The table also lists the ratio of peak material volume concentration to the background level. These peak/background ratios were inconsistent; no trends were observed.

Run	Process	Max/Background	Number of Samples to Return to Background	
	Pipe Discharge			
2	Liquid discharge 35 gpm (0.0022 m ³ /s)	11.5	> 64	
3 – 4	Liquid discharge 105 and 179 gpm $(0.00662 \text{ and } 0.0.0113 \text{ m}^3/\text{s})$	34	> 35	
11	Slurry discharge 43 gpm (0.00271 m ³ /s)	40	> 28	
12	Slurry discharge 125 gpm (0.00789 m ³ /s)	4.6	68	
	Transfer Jet			
5	Liquid discharge 32 gpm (0.00202 m ³ /s)	14	> 12	
6	Liquid discharge 57 gpm (0.00360 m ³ /s)	2.5	51	
10	Slurry discharge 23 gpm (0.00145 m ³ /s)	8.6	99	
	Cutting Jet			
7	Jet on steel at low pressure	4.3	51	
8	Jet on simulant at low pressure	97	>78	
9	Jet on simulant at high pressure	10	62	

Table 6.1.	Sampling cycles for OPC data to return to background and maximum/background						
concentration ratios							

6.2 Particle Size Results

The particle count data from the OPC was converted to a volume (equivalent to mass for homogenous particles) within each size bracket. As shown in Figure 6.2, the data were reasonably well represented by a unimodal (single peak) distribution; so they were fitted to log-normal distributions. In most cases, the secondary fine particle mode had only a minor contribution to the particle volume distribution. The distribution parameters are the geometric mean diameter^{a)} (GMD) and the geometric standard deviation^{b)} (GSD). The results are tabulated in Table 6.2 for each run and for the time intervals that represented the background and peak concentrations during the operation. Even thou,gh data in Table 6.1 showed that the aerosol concentration was markedly elevated during the operations most of the time, the GMD

^{a)} Half of the total aerosol volume is associated with particles larger than the GMD, and the other half with smaller particles. The units are micrometers (μ m).

^{b)} The GSD represents the spread of the data. A GSD of one means that all particles are of the same size.

remained about the same for the 2-in. and 9/16-in. (1.43 cm) nozzles. When the high pressure spray was used, the <u>GMD decreased</u>.

Figure 6.2. Run 2 particle size plot during the background measurement^{a)}

Dun	Drogoss	Backg	round	Peak		
Kull	Trocess	GMD µm	GSD µm	GMD µm	GSD μm	
1	Background	6.0	1.7	N.A.	N.A.	
	Pipe Discharge					
2	Liquid discharge 35 gpm (0.0022 m ³ /s)	6.6	1.8	7.5	1.4	
3–4	Liquid discharge 105 and 179 gpm (0.00662 and 0.0.0113 m ³ /s)	6.2	1.7	6.8	1.4	
11	Slurry discharge 43 gpm (0.00271 m ³ /s)	4.0	2.3	6.7	1.4	
12	Slurry discharge 125 gpm (0.00789 m ³ /s)	6.9	1.6	6.4	1.4	
	Transfer Jet					
5	Liquid discharge 32 gpm (0.00202 m ³ /s)	7.4	1.7	7.0	1.4	
6	Liquid discharge 57 gpm (0.00360 m ³ /s)	8.0	1.4	7.2	1.3	
10	Slurry discharge 23 gpm (0.00145 m ³ /s)	6.5	1.6	5.0	1.6	
	Cutting Jet					
7	Jet on steel at low pressure	8.1	1.4	7.1	1.3	
8	Jet on simulant at low pressure	5.4	1.6	4.0	1.5	
9	Jet on simulant at high pressure	6.4	1.4	3.2	1.5	
	Average	6.5	1.7	6.1	1.4	

 Table 6.2.
 Particle size data from the OPC

^{a)} On the y-axis, the arbitrary mass units are divided by the difference in logarithms of the boundaries of each particle size range to remove the bias in the histogram from the logarithmically irregular size brackets.

6.3 Release Fractions

Appendix C shows the detailed calculations of release fractions. In instances where samples were damaged or flow was lost, the results for those samples were disregarded. Observations and notations are provided with the calculations.

The release fraction results are tabulated in Table 6.3 by location in the ¹/₄-scale tank and for each process simulated. The fluorometry based release fractions represent the release of the liquid fraction of the process slurry. There is an entry in the table for each air sampler unless the sampler pump malfunctioned and no data was obtained for that sampler.

		Release Fraction Data fro Water and from the	om Fluorescent Tracer • Dry Solids (in parenth	in Process heses) Near process stream 2.2 x 10 ⁻⁶ 2.1 x 10 ⁻⁶ 4.5 x 10 ⁻⁶ 4.1 x 10 ⁻⁶
Run	Process	Tank top	Tank Wall	Near process stream
	Pipe Discharge			
2	Liquid discharge 35 gpm (0.0022 m ³ /s)	0.69, 0.80, and 0.93 x 10 ⁻⁶	0.55 and $1.2 \ge 10^{-6}$	2.2 x 10 ⁻⁶
11	Slurry discharge 43 gpm (0.00271 m ³ /s)	0.43, 1.5 and 1.7 x 10^{-6}	2.1 and 2.4 x 10^{-6}	2.1 x 10 ⁻⁶
12	Slurry discharge 125 gpm (0.00789 m ³ /s)	3.1, 3.7 and 4.1 x 10^{-6}	5.0 x 10 ⁻⁶	4.5 x 10 ⁻⁶
	Transfer Jet			
5	Liquid discharge 32 gpm (0.00202 m ³ /s)	0.48 and 3.2×10^{-6}	$3.6 \text{ and } 5.0 \ge 10^{-6}$	4.1 x 10 ⁻⁶
10	Slurry discharge 23 gpm (0.00145 m ³ /s)	1.8, 2.1 and 2.1 x 10^{-6}	2.5 and 2.7 x 10^{-6} (3.5 and 6.8 x 10^{-6})	1.6 x 10 ⁻⁵ (4.4 x 10 ⁻⁵)
	Cutting Jet			
7	Jet on steel at low pressure	1.0 and 5.6 x 10^{-4}	$0.12 \text{ and } 6.3 \text{ x } 10^{-4}$	$4.0 \ge 10^{-4}$
8	Jet on simulant at low pressure	5.5 and 7.7 x 10^{-5}	N.A.	$\frac{1.4 \times 10^{-4}}{(1.3 \times 10^{-4})}$
9	Jet on simulant at high pressure	6.0, 6.0 and 8.1 x 10 ⁻⁵ (6.6, 6.6, and 7.9 x 10 ⁻⁵)	8.8 and 9.1x 10 ⁻⁵ (3.9 and 3.9 x 10 ⁻⁵)	8.0 x 10 ⁻⁵

As noted in section 5.3.1, the results for Run 7 are for the cutting jet on bare steel. The cutting jet may be used to scarify solid waste that is otherwise not sufficiently mobilized by other dislodging mechanisms. If the water-based cutting jet actually strikes bare steel, it is not striking waste, and waste is not being released at that particular time. Consequently, a measured release fraction from this run is not actually applicable to waste retrieval. The results from Runs 8 and 9 are the appropriate release fractions to use for this purpose. The process simulation only contained a solids component for Runs 8 through 12.

Solids release fractions were obtained for only a third of the possible instances (runs X positions), because the measurement method was not as sensitive as it was for the liquid fraction. Agreement between the liquid and solid release fractions was variable but always within a factor of 3. However, in the one case (Run 9) where both release fractions were measurable at the top of the tank, the results showed excellent agreement among the three samplers.

7.0 References

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Appendix A

Air Sampling Procedure for Suspension Experiment

Test Preparation

- 1. Obtain and characterize simulants
- 2. Obtain waste process simulators
- 3. Obtain 0.01 mg capable balance and verify that it is in calibration
- 4. Obtain fluorometer and its calibration data
- 5. Obtain fluorometer procedure
- 6. Obtain sample filter leaching solutions (distilled water) and containers
- 7. Obtain air sample filter holders, filters, cassettes, caps
- 8. Obtain air sampler calibrator
- 9. Obtain optical particle counter interfaced with laptop and data logging software. Record particle data in differential mode.
- 10. Obtain dust monitor(s), interface with laptop and data logging software
- 11. Calculate quantity of dye (fluorescein) needed for makeup water
- 12 Obtain determined quantity of dye (dry or solution) sufficient for each batch of makeup water
- 13. Determine process simulator operating parameters
 - a. Pressure
 - b. Run time
 - c. Configuration
 - d. Sampler run strategy or test instructions
- 14. Prepare data sheets
- 15. Calibrate fluorometer
- 16. Calibrate air sampling pumps and record on data sheets.
 - a. Calibrate the sampling pump using a representative loaded IOM Sampler^{a)} in-line.
 - b. Using flexible tubing, connect the outlet of the IOM Calibration Adapter (Figure A.1) to a flow meter.
 - c. Load a representative filter into an IOM cassette. Insert the cassette into a clean IOM Sampler body. Attach the Calibration Adapter to the IOM by raising the adapter's flat plate (held by springs on both ends) sufficiently to place the IOM front cover face down onto the foam pad.
 - d. Ensure that the IOM front cover is positioned centrally.
 - e. Connect the IOM Sampler outlet to the inlet of a sampling pump.
 - f. Adjust the flow rate to 2 L/min. See calibrator and sampling pump operating instructions for additional information.
 - g. Disconnect the representative sampler from the pump and calibrator and set aside for flow verification after sampling.

^{a)} SKC Incorporated, IOM Inhalable Particle Sampler, Eighty Four, Pennsylvania. This sampler was developed at the Institute of Occupational Medicine, Edinburgh, UK by Mark and Vincent (1986).

Figure A.1. Calibration adaptor

Instructions for Each Test

- 1. Prepare air sampling cassettes and record tare weights on data sheet for the test. Use 25 mm membrane filters. Place in Ziploc bag. Prepare enough units for sampling plus two for field blanks.
 - a. Handling: wear gloves when handling cassettes and use tweezers when working with filters or foams to prevent the transfer of moisture, dust, or other contaminants onto the sampling media.
 - b. Cleaning sampler
 - i. Disassemble the IOM Sampler (Figure A.2). The foam disc will not be used.

Figure A.2. IOM sampler

- Place parts in an ultrasonic cleaner with water and a wetting agent such as soap.
 IOM components may also be cleaned with a solvent such as isopropyl alcohol.
 O-rings should be cleaned separately using water only.
- iii. Clean the components using a lint-free paper or cloth, or a soft lint-free brush. Allow components to dry completely
- c. Loading a filter into the IOM cassette
 - i. Use gentle pressure to separate the two halves of the cassette (Figure A.3)
 - ii. Place a filter into the cassette rear (on the support grid). Snap the cassette front into the cassette rear, ensuring a tight fit.

Figure A.3. Cassette assembly

- d. General weighing guidelines
 - i. Pre- and post-weigh filter and cassette as a single unit. Before pre- and postweighing, wipe the external surface with a clean lint-free paper or cloth, or a soft lint-free brush.
 - ii. Field blanks can be used to correct weights when using plastic cassettes at low filter loadings
 - iii. The IOM cassette assembled with filter or the filter alone should be weighed on a five-figure forced balance. Allow the cassette three-and-half minutes to stabilize before taking a reading. The same balance should be used for both pre- and post-weighing procedures.
- e. Pre-weighing
 - i. Equilibrate cassette and filter at room temperature and humidity for 30 minutes.
 - ii. Wipe the external surface of a loaded IOM cassette with a clean lint-free paper, cloth, or soft brush.
 - iii. Pre-weigh the loaded IOM cassette as a single unit (Figure A.4).
 - iv. Note the result and reference it to the cassette or complete IOM sampler with a number or letter.

- v. Note: The cassette cover (Figure A.4) can be included in pre- and post weighing as part of the complete cassette if required, but must be referenced to the cassette used.
- f. Assemble the cassette in the IOM sampler as shown in Figure A.5. Place in Ziploc® bag for transport to the test site. The large black cap can also be installed over the assembly for transport and run setup.

Figure A.4. Weighing cassette

Figure A.5. Cassette assembly

- 2. Add predetermined fluorescent dye to simulant or process makeup water
- 3. Mix process makeup water.
- 4. Position process simulator and waste simulant in tank.
- 5. Position air monitoring instrumentation and data logging computers.
- 6. Enter new logging file names if needed.
- Collect background data for dust monitor and the particle counter for 30 minutes. Run one filter sample for 30 min as a background. Retrieve air sample cassette. Place in Ziploc® bag for transport to lab.
- 8. Replace background air filter cassette with newly loaded and tared cassette or sampler. Remove large black (if used) and red caps from other samplers.
- 9. Collect a sample of the process water (20 50 ml).
- 10. Note height of process water in tank.
- 11. Restart dust monitor, particle counter, air filters
- 12. Start process simulation per predetermined parameters and record actual operating parameters
- 13. Observe logging of dust monitor and particle counter data, air sampler flow rates.
- 14. Terminate process simulation.
- 15. Close data log files and record file names.

- 16. Record height of process water in tank.
- 17. Cap and retrieve air sample cassettes or complete IOM samplers. Place in individual Ziploc® bags for transport to lab.

Analysis of Air Samples

- 1. Remove sample cassettes from IOM samplers.
- 2. Treat field blanks the same as the actual samples.
- 3. Dry cassettes in oven at 90 °F (32° C)for 30 60 min. Note if cassettes were capped or not.
- 4. Allow cassettes to equilibrate at room temperature and humidity for 30 minutes.
- 5. Reweigh air sample cassettes and record results on the data sheets.
- 6. Include three process blank solution samples with each analysis batch for the remainder of the steps.
- 7. Remove sample filters and place in 50 ml beakers or other containers, collection side up.
- 8. Tare weigh the containers with the filters.
- 9. Put about 20 ml of distilled water into the container with the filters
- 10. Agitate containers occasionally for one hour
- 11. Reweigh the containers to determine amount of water added.
- 12. Filter and transfer solution into fluorometer cuvette, about ³/₄ full
- 13. Analyze each cuvette in Turner fluorometer.^{a)} Repeat three times, rotating cuvette between readings. Record gain setting and readings. Convert readings to mass using calibration data.
- 14. Clean and dry non-disposable containers, IOM samplers, and air filter cassettes

^{a)} Turner Model 450 Digital Fluorometer, currently available from Barnstead International, Dubuque, Iowa.

Appendix B

Optical Particle Counter Measured Concentration as a Function of Time

Optical Particle Counter Measured Concentration as a Function of Time

Figure B.1. Run 1 Optical Particle Counter Background Concentration

Figure B.2. Run 2 Optical Particle Counter concentration pipe discharge with water

Figure B.3. Run 3-4 Optical Particle Counter concentration for pipe discharge with water at high flow rate

Figure B.4. Run 5 Optical Particle Counter concentration for transfer jet with water

Figure B.5. Run 6 Optical Particle Counter concentration for transfer jet with Water

Figure B.6. Run 7 Optical Particle Counter concentration for cutting jet impacting steel plate at low pressure

Figure B.7. Run 8 Optical Particle Counter concentration for cutting jet impacting simulant at low pressure

Figure B.8. Run 9 Optical Particle Counter concentration for cutting jet impacting simulant at high pressure

Figure B.9. Run 10 Optical Particle Counter Concentration for transfer jet slurry discharge

Figure B.10. Run 11 Optical Particle Counter concentration for slurry discharge from a pipe at low flow rate Note the elevated concentration caused by setup activities prior to the simulated operations.

Run 12 OPC Particle Volume

Figure B.11. Run 12 Optical Particle Counter concentration for slurry transfer at high flow rate Note the elevated concentration caused by setup activities prior to the simulated operations.

Appendix C

Release Fraction Calculations

Release Fraction Calculations

Table C.1. Run 1 Background

Background, No Process

Let blank value float to view variability and drift

Concluded that it would be good practice to monitor blank in between samples or often.

									Sample	
	Solution	Gain	FL 1	FL2	FL3	Average	Concentrat	Mass	Volume	
Sample	g						g/g	g	L	Comments
										picked up some
										fluorescent
1A1	20.11	50	616	625	624	621.7	2.18E-08	4.39E-07	45	contamination
1A2	20.03	50	-27	-29	-29	-28.3	1.70E-10	3.41E-09	45	
1A3	20.01	50	-26	-29	-30	-28.3	1.70E-10	3.41E-09	45	
1A4	20.00	50	-33	-26	-27	-28.7	1.59E-10	3.18E-09	45	
1A5	20.00	50	-29	-28	-28	-28.3	1.70E-10	3.40E-09	45	
1A6	20.00	50	-17	-29	-30	-25.3	2.70E-10	5.40E-09	22.5	
1A7	20.26	50	-31	27	-26	-10.0	7.81E-10	1.58E-08		
1A8	19.99	50	-13	-16	-18	-15.7	5.92E-10	1.18E-08		
9 BG	19.99	50	-20	-17	-25	-20.7	4.26E-10	8.51E-09		
PB1	20.00	50	-24	-25	-25	-24.7	2.92E-10	5.85E-09		
PB2	19.98	50	-4	-23	23	-1.3	1.07E-09	2.14E-08		
PB3	19.94	50	-15	-15	-12	-14.0	6.48E-10	1.29E-08		
blk 2		50	-24	-26	-24	-24.7	2.92E-10			
blk 3		50	-42	-46	-42	-43.3	-3.29E-10			
blk 4		50	-33	-32	-32	-32.3	3.70E-11			
Run 1 S	tandards						Std. g/g			
	Avg blk	50				-33.4	0			
	Soln 4	50	130	120	137	129.0	5.41E-09			
						slope	3 33E-11			
						intercept	1.11E-09			
Additio	nal readino	rs								
1 10010101	hlk 5	,- 10	0	0	-1	-03				
	Soln 6	10	168	169	170	169.0				

Table C.2	Run 2 Pipe transfer tests	s with water at low flow rat	te

2 inch tank transfer

35 gpm	Tank Volume	81530 liters
	Dye	275.4 g

									Sample	Release	
Sample	Solution	Gain	FL 1	FL2	FL3	Average	Concentration	Mass	Volume	Fraction	Comments
-	g						g/g	g	L		
2B1	20.06	10	41	42	42	41.7	1.13E-08	2.26E-07	97.5	6.86E-07	•
2B2	20.05	10	49	49	50	49.3	1.31E-08	2.62E-07	97.5	7.95E-07	
2B3	19.99	10	59	60	59	59.3	1.54E-08	3.07E-07	97.5	9.34E-07	
2B4	19.99	10	77	79	77	77.7	1.97E-08	3.93E-07	97.5	1.19E-06	
2B5	20.00	10	32	32	32	32.0	9.01E-09	1.80E-07	97.5	5.47E-07	
2B6	20.01	10	71	71	72	71.3	1.82E-08	3.64E-07	48.75	2.21E-06	
2B7	20.02	10	0	0	0	0.0					
2B8	20.24	10	0	-1	0	-0.3					
2B9 BG	20.04	10	0	0	0	0.0	1.56E-09	3.13E-08	49.5	1.87E-07	
2PB1	20.03	10	0	0	0	0.0					
2PB2	20.02	10	0	0	-1	-0.3					
2PB3	20.03	10	0	0	0	0.0					
									Mean	1.06E-06	
blk 2		10	0	0	0	0.0			S.D.	6.04E-07	
blk 3		10	0	3	0	1.0			Rel S.D.	57%	
blk 4		10	-2	0	0	-0.7					
Run 2 S	tandards						Std. g/g				
	Avg. Blan	k				0.1	0				
	Soln 4	10	19	19	22	20.0	5.41E-09				
	Soln 6	10	149	152	150	150.3	3.92E-08				
	Soln 7	10	1669	1666	1671	1668.7	3.90E-07				
						slope	2.3292E-10				
						intercept	1.5612E-09				

Table C.3. Run 5 Transfer jet tests with water

9/16 nozzle

Use results from GAIN 10, except Gain 50 for sample Tank Volume 81530 liters Gain 50 regression from Run 8 Dye 308.27 g

G 1	G 1	a ·	FF 1						Sample	Release	
Sample	Solution	Gain	FL I	FL2	FL3	Average	Concentration	Mass	Volume	Fraction	Comments
5 A 1	<u> </u>	10	2	2	1	2.0	g/g	g	L		··· 1 11 1
SAI	20.13	10	-3	-2	-1	-2.0	< 0 0 E 00	1.005.07		0.015.04	omitted, problem sample
5A2	20.24	10	262	260	259	260.3	6.83E-08	1.38E-06	114	3.21E-06	
5A3	20.31	10	36	36	36	36.0	1.03E-08	2.09E-07	114	4.85E-07	
5A4	20.20	10	294	295	294	294.3	7.71E-08	1.56E-06	114	3.61E-06	
5A5	20.20	10	408	408	407	407.7	1.06E-07	2.15E-06	114	4.99E-06	
											Used Gain 50 standards
5A6	20.17	50	800	795	791	795.3	3.97E-08	8.00E-07	51	4.15E-06	from Run 8
5A7	20.22	10	-3	-2	-4	-3.0					
5A8	20.14	10	10	9	9	9.3			Mean	3.29E-06	
5B9 BG	Not used,	was ar	nother	backs	ground	that day			S.D.	1.70E-06	
5PB1	21.14	10	0	0	0	0.0			Rel S.D.	52%	
5PB2	20.13	10	-5	-2	-4	-3.7					
5PB3	20.11	10	0	0		0.0					
blk 2		10	0	0	0	0.0					
Run 5 S	tandards						Std. g/g				
	Avg BLK	+ Proc	cess B	lanks		-0.9	0				
	Soln 4	10	17	17	17	17.0	5.41E-09				
	Soln 6	10	145	144	145	144.7	3.92E-08				
	Soln 7	10	1508	1506	1502	1505.3	3.90E-07				
						slope	2.58E-10				
						intercept	9.94E-10				
Extra re	adings					•					
blk 3	U	50	0	2	0	0.7					
blk 4		50	1	-2	-1	-0.7					
5A6	20.17	10	158			158.0	incomplete read	lings, use	Gain 50 r	eadings	

Table C.4.	Run 7 Cutting jet tests with water on steel at low pr	essure
	real , catching jet tests while water on steel at 10 % pr	

High pressure jet on steel -- 300 psig

Tank Volume	81530 liters
Dye	15.38 g

									Sample	Release	
Sample	Solution	Gain	FL 1	FL2	FL3	Average Co	oncentration	Mass	Volume	Fraction	Comments
	g					g/s	g	g	L		
7B1	21.07	10	3	4	3	3.3	1.52E-09	3.20E-08	61.5	2.76E-06	omitted fr. Avg.
7B2	20.05	10	1257	1257	1247	1253.7	3.22E-07	6.45E-06	61.5	5.56E-04	
7B3	20.06	10	213	214	213	213.3	5.53E-08	1.11E-06	61.5	9.56E-05	
7B4	20.08	10	1422	1422	1421	1421.7	3.65E-07	7.32E-06	61.5	6.31E-04	
7B5	20.04	10	25	26	27	26.0	7.32E-09	1.47E-07	61.5	1.26E-05	
7B6	20.02	10	451	453	451	451.7	1.16E-07	2.33E-06	30.75	4.01E-04	
7B7	21.79	10	0	0	0	0.0					
7B8	21.08	10	0	0	0	0.0			Mean	3.39E-04	
7B9 BG	Not used, v	vas ano	ther ba	ckgrou	nd that	day			S.D.	2.75E-04	
7PB1	21.15	10	0	2	2	1.3			Rel S.D.	81%	
7PB2	21.12	10	0	0	0	0.0					
7PB3	21.09	10	0	0	0	0.0					
blk 2		10	0	0	0	0.0					
Run 7 St	andards						Std. g/g				
	Avg BLK -	+ Proce	ss Blar	ıks		0.3	0				
	Soln 4	10	17	17	17	17.0	5.41E-09				
	Soln 6	10	149	149	149	149.0	3.92E-08				
	Soln 7	10	1522	1520	1522	1521.3	3.90E-07				
					5	slope	2.56E-10				
					i	intercept	6.65E-10				

Table C.5. Run 8 Cutting jet tests on	n simulant at low	pressure
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Tank Volume

81530 liters

High pressure jet on simulant -- 300 psig

							Dye	11.6	g		
Sample	Solution g	Gain	FL 1	FL2	FL3	Average	Concentration g/g	Mass g	Sample Volume L	Release Fraction	Comments
8A1	21.23	50	32	28	29	29.7	1.79E-09	3.80685E-08	57	4.69E-06	omitted, results similar to those that did not run
8A2	21.23	50	424	423	424	423.7	2.09E-08	4.43648E-07	57	5.47E-05	
8A3	21.33	50	596	592	592	593.3	2.91E-08	6.21345E-07	57	7.66E-05	
8A4	21.30	50	32	39	38	36.3	2.12E-09	4.50846E-08			did not run, omitted
8A5	21.18	50	15	17	17	16.3	1.15E-09	2.42976E-08			did not run, omitted
8A6	21.22	50	543	544	543	543.3	2.67E-08	5.66443E-07	28.5	1.40E-04	
8A7	21.16	50	-4	-2	-3	-3.0					
8A8	20.70	50	0	0	-4	-1.3					
8B9 BG	21.11	50	-4	-3	-12	-6.3	4.80E-11	1.01219E-09	45		
8PB1	21.11	50	-3	-4	-2	-3.0					
8PB2	21.12	50	-1	-2	-6	-3.0					
8PB3	20.13	50	1	0	0	0.3			Mean	9.03E-05	
									S.D. Rel S.D.	4.41E-05 49%	
blk2		50	0	0	2	0.7				.,,,	
Run 8 St	andards						Std. g/g				
	Avg BLK -	+ Proce	ess Bla	anks		-1.3	0				
	Soln 4	50	99	96	97	97.3	5.41E-09				
	Soln 6	50	803	801	802	802.0	3.92E-08				
					:	slope	4.849E-11				
					i	intercept	3.550E-10				
Extra Da	ita					g/g					

						00
8A2	10	71	72	71	71.3	
8A6	10	113	114	113	113.3	
blk 1	10	0	0	0	0.0	
Soln 7	10	1653	1650	1647	1650.0	3.90E-07

Table C.6.	Run 9	Cutting	iet tests	on simula	ant at high	pressure
		C arrang	Jee	011 0111010	me we mon	p1000010

High pressure jet ON clay -- 1200 psig

	-	-		-	_	_	Tank Volume	81530	liters		
							Dye	22.25	g		
									Sample	Release	
Sample	Solution	Gain	FL 1	FL2	FL3	Average	Concentration	Mass	Volume	Fraction	Comments
	g						g/g	g	L		
9B1	21.09	50	814	811	815	813.3	4.06E-08	8.55871E-07	51	6.15E-05	
9B2	21.24	50	1066	1067	1062	1065.0	5.30E-08	1.12641E-06	51	8.09E-05	
9B3	21.30	50	732	732	729	731.0	3.65E-08	7.77634E-07	51	5.59E-05	
9B4	21.30	50	1164	1160	1143	1155.7	5.75E-08	1.22513E-06	51	8.80E-05	
9B5	21.27	50	1195	1191	1187	1191.0	5.93E-08	1.26059E-06	51	9.06E-05	
9B6	21.10	50	531	525	524	526.7	2.64E-08	5.57036E-07	25.5	8.00E-05	
9B7	20.19	50	0	0	5	1.7					
9B8	20.09	50	-1	3	-2	0.0			Mean	6.96E-05	
9B9 BG	Not used,	was a	nother	backg	ground th	nat day			S.D.	1.28E-05	
9PB1	21.15	50	-2	0	-2	-1.3			Rel S.D.	18%	
9PB2	21.11	50	0	-1	0	-0.3					
9PB3	21.17	50	0	1	1	0.7					
			0		0						
biki		50	0	1	0	0.3					
Run 9 S	tandards						Std. g/g				
	Avg BLK	+ Pro	cess E	slanks		-0.2	0				
	Soln 4	50	93	95	96	94.7	5.41E-09				
	Soln 6	50	785	787	787	786.3	3.92E-08				
						slope	4.947E-11				
						intercept	3.444E-10				
Extra D	ata		_			g/g					
8A2	10	71	72	71	71.3						
8A6	10	113	114	113	113.3						
blk 2	10	0	0	0	0.0						
Soln 7	10	1653	1650	1647	1650.0	3.90E-07					

Table C.7.	Run 10	Transfer	jet tests	with slurry	^v discharge
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9/16 nozzle with clay

Tank Volume81530 litersDye295.97 g

									Sample	Release	
Sample	Solution	Gain	FL 1	FL2	FL3	Average	Concentration	Mass	Volume	Fraction	Comments
	g						g/g	g	L		
10A1	21.06	50	604	606	604	604.7	3.19E-08	6.71E-07	103.5	1.79E-06	
10A2	21.15	50	693	696	693	694.0	3.66E-08	7.73E-07	103.5	2.06E-06	
10A3	21.42	50	709	711	705	708.3	3.73E-08	7.99E-07	103.5	2.13E-06	
10A4	20.50	50	906	902	903	903.7	4.76E-08	9.75E-07	100.5	2.67E-06	
10A5	20.07	50	908	898	898	901.3	4.74E-08	9.52E-07	103.5	2.53E-06	
											clay
											deposit on
											sampler
10A6	42.25	50	1340	1330	1322	1330.7	6.99E-08	2.96E-06	51.75	1.57E-05	inlet
10A7	20.55	50	-5	-5	0	-3.3					
10A8	21.08	50	-3	-2	-4	-3.0			Mean	4.48E-06	
10AG9	Not used,	was a	nother	backg	ground	that day			S.D.	5.52E-06	
10PB1	20.99	50	-7	0	-3	-3.3			Rel S.D.	123%	
10PB2	20.70	50	0	-2	-1	-1.0					
10PB3	21.01	50	1	0	0	0.3					
blk 1		50	-1	0	0	-0.3					
blk 2		50	-1	-2	-1	-1.3					
Run 10	Standards						Std. g/g				
	Avg BLK					-0.8	0				
	Soln 4	50	96	98	97	97.0	5.41E-09				
	Soln 6	50	746	747	741	744.7	3.92E-08				
						-1	5 04E 11				
						siope	J.24E-11				
						intercept	1./4E-10				

Table C.	8. Run	11	Pipe	discharge	tests using	g slurry	/ discharge	at high f	flow rate
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2 in. nozzle 30 gpm

Tank Volume81530 litersDye271.79 g

									Sample	Release	
Sample	Solution	Gain	FL 1	FL2	FL3	Average Cond	centration	Mass	Volume	Fraction	Comments
	g					g/g		g	L		
11B1	21.19	50	103	107	105	105.0	5.75E-09	1.22E-07	84	4.35E-07	
11B2	20.57	50	383	384	383	383.3	2.06E-08	4.23E-07	84	1.51E-06	
11B3	21.11	50	412	409	410	410.3	2.20E-08	4.64E-07	84	1.66E-06	
11B4	21.07	50	526	526	525	525.7	2.81E-08	5.93E-07	84	2.12E-06	
11B5	21.07	50	381	396	395	390.7	2.10E-08	4.42E-07	54	2.45E-06	
11B6	21.22	50	254	253	251	252.7	1.36E-08	2.89E-07	42	2.06E-06	
11 B 7	20.05	50	3	0	0	1.0					
11B8	21.05	50	2	1	0	1.0					
11BG9	20.15	50	0	0	3	1.0	2.18E-10	4.39E-09	60	2.20E-08	
11PB1	19.99	50	0	0	3	1.0					
11PB2	20.15	50	-1	4	0	1.0					
11PB3	20.01	50	2	0	0	0.7			Mean	1.71E-06	
									S.D.	7.08E-07	
blk 1		50	0	-2	1	-0.3			Rel S.D.	42%	
Run 11	Standards						Std. g/g				
	Avg BLK	+ Pro	cess b	lanks		0.6	0				
	Soln 4	50	95	90	98	94.3	5.41E-09				
	Soln 6	50	743	731	728	734.0	3.92E-08				
						slope	5.32E-11				
						intercept	1.65E-10				

Table C.9.	Run 12 Pi	pe transfer	tests of slurry	discharge	at high flow rate
			2		

2 in. nozzle 126 gpm

Tank Volume 81530 liters Dye 275.73 g

									Sample	Release	
Sample	Solution	Gain	FL 1	FL2	FL3	Average Cond	centration	Mass	Volume	Fraction	Comments
	g					g/g		g	L		
12B1	20.10	50	507	512	507	508.7	2.73E-08	5.49E-07	52.5	3.09E-06	
12B2	21.16	50	582	578	577	579.0	3.10E-08	6.57E-07	52.5	3.70E-06	
12B3	21.23	50	646	646	643	645.0	3.46E-08	7.34E-07	52.5	4.13E-06	
12B4	20.11	50	823	820	817	820.0	4.39E-08	8.82E-07	52.5	4.97E-06	
											omitted
											from
12B5	21.18	50	7	7	6	6.7	6.24E-10	1.32E-08	52.5	7.44E-08	Average
12B6	21.01	50	353	354	358	355.0	1.91E-08	4.02E-07	26.25	4.53E-06	
12B7	20.04	50	0	1	0	0.3					
12B8	21.09	50	0	2	-5	-1.0			Mean	4.08E-06	
12BG9	Not used,	was a	nother	back	groun	d that day			S.D.	7.27E-07	
12PB1	21.10	50	-2	-7	0	-3.0			Rel S.D.	18%	
12PB2	20.09	50	-2	-2	-1	-1.7					
12PB3	20.00	50	0	0	-1	-0.3					
blk 1		50	1	-2	0	-0.3					
blk 2		50	0	-1	-2	-1.0					
Run 12	Standards						Std. g/g				
	Avg BLK					-0.7	0				
	Soln 4	50	89	94	92	91.7	5.41E-09				
	Soln 6	50	732	730	737	733.0	3.92E-08				
						slope	5.32E-11				
						intercept	2.70E-10				

Table C.10. Process water samples

Process water samples

Sample	D.I water	Solution	Gain	GL1	FL2	FL3	Average	Solution	Process Water	Process Water Used In Run	Dye Exposed in Run	Comment
D 2	<u> </u>	20.0114	1	500	596	570	5047	<u>g/g</u>	<u>g/g</u>	<u>g</u>	g 275.40	
Run Z	30.4955	30.9114	1	389	380	5/9	584.7	1.29E-06	9.62E-05	2.86E+06	275.40	
Run 5	31.2173	31./116	1	762	762	760	761.3	1.68E-06	1.08E-04	2.86E+06	308.27	
Run 7	30.3107	30.6687	1	459	458	454	457.0	1.02E-06	8.71E-05	1.77E+05	15.38	
Run 8	29.9810	30.2078	1	261	260	260	260.3	5.89E-07	7.85E-05	1.48E+05	11.60	
Run 9	29.9446	30.2066	1	275	276	276	275.7	6.23E-07	7.18E-05	3.10E+05	22.25	
Run 10	29.0415	29.6414	1	953	952	950	951.7	2.09E-06	1.03E-04	2.86E+06	295.97	Probably contained some clay,
Run 11	29.0411	29.3979	1	519	520	519	519.3	1.15E-06	9.50E-05	2.86E+06	271.79	although centrifuged before
Run 12	29.7669	30.1634	1	573	572	570	571.7	1.27E-06	9.64E-05	2.86E+06	275.73	analysis
10PB1			1	0	0	0	0.0					
10PB2			1	0	0	0	0.0		9.08E-05	Mean		
10PB3			1	0	0	0	0.0		1.42E-05	S.D.		
City wate	r		1	0	0	0	0.0		16%	Rel S.D.		
blk 1			1	0	0	0	0.0					
	-	D-4-1. 64						C(1 - /-				
	1	Satch Star	idards				0.0	Sta. g/g				
	1	Avg BLK 4	- city v	vater -	+ proc	ess bl	0.0	0				
	S	Soln 3	1	932	929	927	929.3	2.04E-06				
	S	Soln 7	1	157	155	156	156.0	3.90E-07				

slope 2.18E-09 intercept 2.30E-08

High pressure jet on simulant 300 psig								
	Tank Vo	olume	81530	liters				
		Clay	4491	g				
Sample	Weight Sample Change Volume mg	L	Release Fraction	Comment				
12B1	0.0	57	0.00E+00	omitted from fluorometry and here				
12B2	0.1	57	3.18E-05					
12B3	0.0	57	0.00E+00	omitted zeroes				
12B4	0.0			did not run, omitted				
12B5	0.0			did not run, omitted				
12B6	0.2	28.5	1.27E-04					
12B7	0.0							
12B8	0.0		7.96E-05	Mean				
12BG9	0.1		6.76E-05	S.D.				
			85%	Rel S.D.				

Table C.11. Run 8 Solids analysis from cutting jet impacting simulant at low pressure

Table C.12. Run 9 Solids analysis from cutting jet impacting simulant at high pressure

High pressure jet on simulant -- 1200 psig Tank Volume 81530 liters

		Clay	11839	g
Sample	Weight Sample Change Volume	F F	Release Fraction	Comment
	mg	L		
12B1	0.5	52.5	6.56E-05	
12B2	0.5	52.5	6.56E-05	
12B3	0.6	52.5	7.87E-05	
12B4	0.3	52.5	3.94E-05	
12B5	0.3	52.5	3.94E-05	
12B6	0.1	26.25	2.62E-05	
12B7	-0.2			
12B8	-0.3		5.25E-05	Mean
12BG9	N.A.		2.03E-05	S.D.
			39%	Rel S.D.

 Table C.13. Run 10 Solids analysis from transfer jet slurry discharge

9/16 nozzle with slurry

	Tank	Volume	81530	liters
		Clay	46040	g
	Weight	Sample	Release	
Sample	Change	Volume	Fraction	Comment
	mg	L		
12B1	0.0	103.5	0.00E+00	omitted zeroes
12B2	-0.1	103.5	-1.71E-06	omitted zeroes
12B3	0.0	103.5	0.00E+00	omitted zeroes
12B4	0.2	100.5	3.52E-06	
12B5	0.4	103.5	6.84E-06	
12B6	1.3	51.75	4.45E-05	clay deposit on sampler inlet
12B7	-0.1			
12B8	-0.1		1.83E-05	Mean
12BG9	N.A.		2.28E-05	S.D.
			124%	Rel S.D.

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