



Total Dissolved Residue (Total Dissolved Solids, TDS) in Aqueous Matrices

Environmental Express

2345A Charleston Regional Parkway

Charleston, SC 29492

800-343-5319

Table of Contents

1. Scope and Application.....	2
2. Summary of Method.....	2
3. Definitions	2
4. Interferences	3
5. Safety	3
6. Equipment and Supplies	4
7. Reagents and Standards	5
8. Sample Collection Preservation and Storage.....	5
9. Quality Control	6
Table 1: 40 CFR part 136.7 Quality Control Requirements	6
10. Procedure	9
11. Data Analysis and Calculations.....	10
12. Method Performance	10
13. Pollution Prevention	11
14. Waste Management	11
15. References	11
16. Tables, Diagrams, Flowcharts, and Validation Data	12
Table A1: Environmental Express TDS Stable-Weigh Tests Blank	13
Table A2: Environmental Express TDS Stable-Weigh Tests 200 mg.....	15
Table A3: Environmental Express TDS Stable-Weigh Tests 100 mg.....	17
Table A4: Environmental Express TDS Stable-Weigh Tests 20 mg.....	19
Table A5: Environmental Express TDS Stable-Weigh Tests 20 mg-Evaporation Dish ..	21
Table A6: StableWeigh and Porcelain Evaporation Dish Precision.....	23
Table A7: StableWeigh Method Blank.....	23

Table of Figures

Figure 1: Initial Demonstration of Capability Calculation	7
Figure 2: LRB Calculation.....	7
Figure 3: LFB Calculation	7
Figure 4: RPD Calculation.....	8
Figure 5: Total Solids Calculation	10
Figure 6: StableWeigh 200 mg Initial Demonstration of Capability	24
Figure 7: StableWeigh 100 mg Initial Demonstration of Capability	25
Figure 8: StableWeigh vs. Porcelain Evaporation Dish Averages	26

1. Scope and Application

- 1.1. This method is based upon Standard Methods 2540 for the determination of Total Dissolved Residue in aqueous sample matrices.
- 1.2. This method is for use in the Environmental Protection Agency's (EPA's) data gathering and monitoring programs under the Clean Water Act, the Resource Conservation and Recovery Act, the Comprehensive Environmental Response, Compensation and Liability Act, and the Safe Drinking Water Act.
- 1.3. This method is used to determine Total Dissolved Residue (Total Dissolved Solids, TDS).
- 1.4. This method is applicable to drinking, surface, and saline waters, domestic and industrial wastes
- 1.5. The concentration range for this method is determined by the minimum and maximum residue mass allowed (2.5 – 750 mg) and the volume of sample needed to achieve the desired residue mass. Samples for TDS have no upper volume limit specified but are limited by the size of the evaporation vessel and the practicality of repeated additions of sample volume.
- 1.6. Each laboratory that uses the method must demonstrate the ability to generate acceptable results using the procedure.

2. Summary of Method

- 2.1. For TDS the entire aliquot of sample is evaporated in a pre-weighed vessel and dried to constant weight at $180 \pm 2^\circ\text{C}$. The mass of the residue is determined by difference in mass of the vessel.

3. Definitions

- 3.1. Laboratory Fortified Blank (LFB) - An aliquot of reagent water or other blank matrix to which known quantities of the method analytes and all the preservation compounds are added. The LFB is processed and analyzed exactly like a sample, and its purpose is to determine whether the methodology is in control, and whether the laboratory is capable of making accurate and precise measurements.
- 3.2. Laboratory Matrix/Duplicate (LM/LMD) also called Matrix / Duplicate (M/D): An aliquot of an environmental sample and a duplicate which are analyzed and its purpose is to determine whether the sample matrix contributes bias to the analytical results.
- 3.3. Laboratory Reagent Blank (LRB) - A volume of reagent water or other blank matrix that is processed exactly as a sample including exposure to all glassware and equipment, that

are used in the analysis batches. The LRB is used to determine if the method analytes or other interferences are present in the laboratory environment, the reagents, or the apparatus.

- 3.4. Minimum Reporting Level (MRL) - The minimum concentration that can be reported by a laboratory as a quantitated value for a method analyte in a sample following analysis. This concentration must not be any lower than the concentration of the lowest calibration standard for that instrument.
- 3.5. Total Dissolved Residue or Total Dissolved Solids is defined as all material that is not volatile at 180°C.
- 3.6. Water Sample: For the purpose of this method, a sample taken from one of the following sources: drinking water, surface water, storm runoff, industrial or domestic wastewater.

4. Interferences

- 4.1. Samples with multiple phases or those that settle into multiple layers present a problem with homogeneity. Ensure that such samples are well mixed during use. Vigorous shaking through inversion will provide a sufficient mixing for most samples. Samples that settle rapidly or resist mixing may require constant stirring with a magnetic stirrer. Use a wide bore pipette in such instances. Draw sample from the midpoint between the wall and the vortex. Avoid using propeller style mixers as these may shear particles resulting in underreported values.
- 4.2. Some samples may form a crust during drying. This crust can prevent some evaporation of water during the drying process. If a water trapping crust continues to be problematic reduce the amount of sample used to minimize the crust being formed or take other steps to allow water to evaporate. Document additional steps being used as well as the precautions that are taken to avoid loss of residue from the dried sample.
- 4.3. Organic compounds such as oil and grease may present problems in obtaining a stable final weight. Such compounds are often visible to the naked eye and will present as a constantly decreasing mass after each drying event. Develop a Quality Plan to positively identify these types of samples. Obtain a value for HEM for such samples, if possible and consult with the final data user how to use that value in conjunction with the data obtained from the analysis.

5. Safety

- 5.1. When working with and around ovens or other evaporation equipment the materials in use will become hot. Use appropriate heat resistant gloves and other necessary PPE to protect the analysts from exposure to heat. Different temperatures may require different PPE to be effective.

- 5.2. Use all standard PPE (gloves, safety glasses, lab coats, etc.) as required by your Laboratory Safety and Chemical Hygiene Plan. Pay particular attention to samples of unusual appearance or odor and request additional information from the sample source as necessary to identify unknown or unexpected hazards.
- 5.3. This method does not address all safety issues associated with its use. The laboratory is responsible for maintaining a safe work environment and a current awareness file of OSHA regulations regarding the safe handling of the chemicals specified in this method. A reference file of safety data sheets (SDS's) must be available to all personnel involved in these analyses.

6. Equipment and Supplies

Note: Brand names, suppliers and part numbers are cited for illustrative purposes only. No endorsement is implied. Equivalent performance may be achieved using equipment and materials other than those specified here, but demonstration of equivalent performance that meets the requirements of this method is the responsibility of the laboratory.

- 6.1. Evaporation Vessel – StableWeigh™ TDS Vessel from Environmental Express, or equivalent. This vessel is not suitable for use in volatile solids determination. In such instances use ceramic, high silica glass, or platinum evaporating dishes of appropriate dimensions. When using the StableWeigh™ product, other accessories are recommended or required for ease of use.
- 6.2. StableWeigh™ modular rack – recommended. Used to transport and hold StableWeigh™ Vessels to, from, and in ovens and desiccators.
- 6.3. StableWeigh™ weighing bracket – recommended. Used to support vessels on the balance.
- 6.4. Static electricity dissipation device. Best results have been achieved using the Mettler Toledo balance deionizer or similar device. Mounted anti-static ionizing cartridges or brushes can be effective if care is taken to ensure the entire vessel is appropriately treated.
- 6.5. Drying oven for use at 103-105°C and 178-182°C.
- 6.6. Dessicator with dessicant– must include some indicator to determine when replacement of the desiccant is necessary. This can be indicating desiccant or other appropriate means.
- 6.7. Analytical balance capable of accurate and precise weighing to 0.1 mg.
- 6.8. Class A measuring devices of appropriate volume to measure necessary amounts of sample.

- 6.9. Glass Fiber Filter Discs – without organic binder, and 1.5 μ m nominal pore size. Unprepared filters are available or use Washed & Dried filters from Environmental Express, or equivalent, to skip the filter preparation steps in the procedure.
- 6.10. Filtration apparatus, StableWeigh™ Filling Station – Use a filter funnel of the appropriate size to hold the volume of sample and diameter of filter being used. The StableWeigh™ Filling Station (single place or six place) will allow the analyst to filter samples directly into the vessel.

7. Reagents and Standards

- 7.1. The various forms of residue are determined based solely on physical characteristics. No specific chemicals or reagents are required for this analysis.
- 7.2. No specific standard is required. The TDS Standard from Environmental Express or equivalent may be used. Each purchased standard may have slightly different target values. Consult the certificate of analysis for the assigned true value of each standard. Alternatively, follow the directions in 9.6.1.2

8. Sample Collection Preservation and Storage

- 8.1. Always refer to the latest guidelines in 40 CFR part 136 for sampling guidelines, preservation guidelines, and holding time. Those instructions supersede any guidance given in this document.
- 8.2. Collect sample in a plastic or glass bottle. A recommended volume is a minimum of 250 mL.
- 8.3. Store samples at $\leq 6^{\circ}\text{C}$ when not in use.
- 8.4. Store samples for no more than 7 days from collection.

9. Quality Control

9.1. Each laboratory that uses this method is required to develop and implement a formal Quality Manual. The minimum requirements of this program are outlined in 40 CFR part 136.7.

Table 1: 40 CFR part 136.7 Quality Control Requirements	
40 CFR 136.7 QC Requirement	Required (Yes/No) and Notes
Demonstration of Capability (DOC),	Initial and Ongoing Demonstration of Capability Required
Method Detection Limit (MDL),	MDL _{Spike} and MDL _{Blank} not required for methods defined test.
Laboratory reagent blank (LRB),	Required daily or with each batch of 20 or fewer samples.
Laboratory fortified blank (LFB), also referred to as a spiked blank, or laboratory control sample (LCS),	LFB is required daily or with each batch of 20 or fewer samples.
Matrix spike (MS), matrix spike duplicate (MSD), or laboratory fortified blank duplicate (LFBD) for suspected difficult matrices,	Duplicates daily or with each batch of 20 or fewer samples.
Internal standard/s, surrogate standard/s (for organic analysis) or tracer (for radiochemistry),	Not required.
Calibration (initial and continuing),	Calibrate balance as per manufacturer and regulatory requirements.
Control charts (or other trend analyses of quality control results),	Required.
Corrective action (root cause analyses),	Required.
Specific frequency of QC checks,	Required.
QC acceptance criteria, and	Required.
Definitions of a batch (preparation and analytical)	Required.

Note: Daily: Each day samples are analyzed in the laboratory.

9.2. Initial Demonstration of Capability (IDC)

- 9.3. Before new analysts run any samples, verify their capability with the method. Run a LFB at least four times. Calculate the standard deviation of the four samples. All 4 LFB values must fall within the LFB recovery limits for the analyst to pass the IDC

$$\text{LFB's initial recovery limits} = \text{Mean} \pm (5.84 \times \text{Standard Deviation})$$

Where:

5.84 = the two-sided Student's t factor for three degrees of freedom.

Figure 1: Initial Demonstration of Capability Calculation

- 9.4. Laboratory Reagent Blank (LRB):
- 9.5. High quality laboratory water that is analyzed as a sample. Analyzed at a frequency of no less than once per sample batch. Develop control chart limits to determine acceptable recovery. Perform Root Cause Analysis as per the laboratory Quality Manual in recovery is unacceptable

$$\text{Experimental Value} = \text{LRB}$$

$$\text{Experimental Value} = \text{LRB Concentration determined experimentally}$$

Figure 2: LRB Calculation

- 9.6. Laboratory Fortified Blank (LFB):
- 9.6.1. LFB can :
- 9.6.1.1. Be purchased from a 3rd party supplier such as Environmental Express. Follow supplier's directions or:
- 9.6.1.2. Prepare in a 1000 ml volumetric flask put 0.2500 g (± 0.0001 g) of Celite and 20.0000 g (± 0.0001 g) of Sodium Chloride. Dilute to volume with DI water and mix well. Transfer 10 ml of the LFB to a StableWeigh vessel and dry. Recovery will be 20.0000 g/L.
- 9.6.2. A maximum recovery range of 80% to 120% is expected. Develop control chart limits to determine acceptable recovery. Perform Root Cause Analysis as per the laboratory Quality Manual in recovery is unacceptable.

$$\left(\frac{\text{Experimental Value}}{\text{Expected Value}} \right) * 100 = \text{Percent Recovery LFB}$$

Experimental Value = LFB Concentration determined experimentally

Expected Value = Known LFB concentration

Figure 3: LFB Calculation

- 9.7. Laboratory Matrix/ Laboratory Matrix Duplicate (LM/LMDD):

9.8. A duplicate set of TS samples are analyzed. LM/LMD are analyzed at a frequency of no less than once per sample batch. A Relative Percent Difference (RPD) will be calculated for each LM/LMD set. Develop control chart limits for RPD to determine acceptable recovery. Perform Root Cause Analysis as per the laboratory Quality Manual in recovery is unacceptable.

$\text{Absolute Value} \left(\frac{\left(\frac{\text{LM-LMD}}{\left(\frac{\text{LM+LMD}}{2} \right)} \right)}{\left(\frac{\text{LM+LMD}}{2} \right)} \right) * 100 = \text{RPD}$
<p>LM = Concentration determined for LM</p> <p>LMD = Concentration determined for LM duplicate</p>

Figure 4: RPD Calculation

9.9. Ongoing Demonstration of Capability

9.10. Samples consisting of the LRB, LFB, and LM/LMD will be run each batch of samples. These results will be charted on control charts to determine ODC for the laboratory. Develop control chart limits to determine acceptable recovery. Perform Root Cause Analysis as per the laboratory Quality Manual in recovery is unacceptable.

9.11. Balance Check. An appropriate check weight should be evaluated at the beginning and end of each weighing sequence. The checks will demonstrate lack of drift in the balance. Reasonable choices would be within the same order of magnitude as the expected mass of the residue plus StableWeigh vessel used for the samples. Develop control chart limits to determine acceptable check weight. Perform Root Cause Analysis as per the laboratory Quality Manual in check weight is unacceptable.

9.12. Sample Batch

9.12.1. A group of samples which behave similarly with respect to the sampling or the testing procedures being employed and which are processed as a unit. For QC purposes, if the number of samples in a group is greater than 20, then each group of 20 samples or less will all be handled as a separate batch. A batch cannot span between laboratory work days (24 hrs.). New batches must be started each laboratory work day.

9.12.2. Sample Batch: Typical sample analysis sequence.

9.12.2.1. Balance Check

9.12.2.2. LRB

9.12.2.3. LFB

- 9.12.2.4. LM
 - 9.12.2.5. LMD
 - 9.12.2.6. Samples
 - 9.12.2.7. Balance Check
 - 9.12.2.8. Other Equipment Checks
- 9.13. Other equipment used in this analysis, such as ovens, balances, and thermometers, must be maintained and checked as required by the laboratory Quality Manual.
- 9.14. Control charts and trend analysis will be performed on all areas identified in the Quality Control section in this method and in the laboratories Quality Manual.

10. Procedure

Note: This method is entirely empirical. Acceptable results can be obtained only by strict adherence to all details.

- 10.1. Select an appropriate number of StableWeigh TDS vessels for the required number of samples and QC. Assign each vessel to a sample and note the vessel tare weight in the appropriate analysis data log. StableWeigh vessels must not be used if the sample aliquot is to be analyzed for volatile solids.
- 10.2. If ProWeigh Washed and Dried filters are used, skip the following preparation steps and proceed to 10.7
- 10.3. Insert a filter into the filtration apparatus and apply vacuum.
- 10.4. Wash the filter with three successive 20 mL portions of reagent water. Continue to apply vacuum to remove all traces of wash water from the filter.
- 10.5. Discard wash water.
- 10.6. Choose a sample size expected to yield between 2.5 and 700 mg of residue. Mix the sample well through appropriate means (repeated inversion, stirring with magnetic stirrer or other) and measure out the appropriate sample volume. If the vessel is not of sufficient size to hold the volume required, successive portions of sample may be added after evaporation.
- 10.7. Place the filter and StableWeigh TDS Vessel in the StableWeigh filling station. Apply vacuum and add the sample to the filter holder. No more than 10 minutes should be required for sample filtration. Larger filter sizes or smaller sample volumes may be necessary to fulfill this time requirement. After filtration is complete wash the filter with three successive 10 mL portions of reagent water to ensure complete transfer of dissolved constituents into the filtrate. Continue suction until all visible water has been removed from the filter.

- 10.8. Remove the StableWeigh TDS vessel from the filling station and transfer it to the oven rack.
- 10.9. Evaporate the sample to dryness using a block digester, oven, steam bath, or other appropriate method. Set the evaporation temperature low enough that the samples do not boil and splatter.
- 10.10. Dry the evaporated sample and vessel in a drying oven at $180 \pm 2^{\circ}\text{C}$ for at least one hour.
- 10.11. Remove the vessels from the oven and place in a dessicator until they have reached balance temperature. Cooling time for StableWeigh vessels can be reduced by transferring the vessels to a cool rack prior to placing in the dessicator.
- 10.12. Weigh each vessel on the balance and record the weight to 0.1 mg.
- 10.13. Repeat steps 11.2.8 - 11.2.10 at least one more time and as many times as necessary to obtain consecutive weights that differ by less than 0.5 mg. QC samples must go through the drying/cooling/weighing cycle as long as other samples require them. Other samples may be removed from the cycle once they have shown weight stability. When using crucibles, keeping the dessicator time constant with each cycle will help to reduce the number of cycles needed to achieve constant weight.

11. Data Analysis and Calculations

- 11.1. Calculations for Total Dissolved Solids

$\frac{(A-B) \times 1000}{C} = \text{Total Dissolved Solids (mg/L)}$ <p>A = Weight of dried sample + StableWeigh (mg)</p> <p>B = Weight of StableWeigh (mg)</p> <p>C = volume of sample (mL)</p>
--

Figure 5: Total Solids Calculation

12. Method Performance

- 12.1. See Standard Methods 2540 for performance data on the original analysis method.
- 12.2. Comparison data validating the use of the StableWeigh vessel in comparison to the crucibles required for TDS are found in Tables/Appendices.

13. Pollution Prevention

- 13.1. The main source of pollution from this method is from unknown samples. Follow your laboratory Waste Management Program for the identification and disposal of potentially hazardous samples.

14. Waste Management

- 14.1. It is the laboratory's responsibility to comply with all federal, state, and local regulations governing waste management, particularly the hazardous waste identification rules and land disposal restrictions, and to protect the air, water, and land by minimizing and controlling all releases from fume hoods and bench operations. Compliance with all sewage discharge permits and regulations is also required.
- 14.2. Use the least amount of sample possible to generate valid results, particularly for TS. This will minimize the amount of waste generated by the analysis.
- 14.3. For further information on waste management, consult "The Waste Management Manual for Laboratory Personnel," and "Less is Better: Laboratory Chemical Management for Waste Reduction," both available from the American Chemical Society's Department of Government Relations and Science Policy, 1155 16th Street N.W., Washington DC, 20036.

15. References

- 15.1. Eaton, A. D., Clesceri, L. S., Baird, R. B., Rice, E. W., American Public Health Association., American Water Works Association., & Water Environment Federation. (2012). *Standard Methods for the Examination of Water and Wastewater, 22nd edition*. Washington, DC: American Public Health Association

16. Tables, Diagrams, Flowcharts, and Validation Data

Table A1: Environmental Express TDS Stable-Weigh Tests Blank

Analyst		Edward F. Askew							
Date		4/14/2016							
Blank									
Sample #	Bag #	Volume (ml)	Initial weight (g)	Final weight 1 (g)	Final weight 2 (g)	Final weight 3 (g)	Two Consecutive Weights Difference (mg)	Final weight used (mg)	Percent Change
1	144	100	3.7783	3.7785	3.7782		-0.3	3.7782	0.00265%
2	145	100	3.7329	3.7330	3.7329		-0.1	3.7329	0.00000%
3	147	100	3.8709	3.8709	3.8708		-0.1	3.8708	0.00258%
4	148	100	3.7953	3.7958	3.7955		-0.3	3.7955	-0.00527%
5	150	100	3.8747	3.8748	3.8745		-0.3	3.8745	0.00516%
6	75	100	3.7540	3.7543	3.7540		-0.3	3.7540	0.00000%
7	58	100	3.7246	3.7249	3.7245		-0.4	3.7245	0.00268%
8	81	100	3.7105	3.7107	3.7105		-0.2	3.7105	0.00000%
9	82	100	3.7503	3.7503	3.7503		0.0	3.7503	0.00000%
10	83	100	3.7500	3.7495	3.7497		0.2	3.7497	0.00800%
11	94	100	3.7312	3.7313	3.7316		0.3	3.7316	-0.01072%
12	86	100	3.7855	3.7854	3.7855		0.1	3.7855	0.00000%
13	77	100	3.9155	3.9153	3.9154		0.1	3.9154	0.00255%
14	85	100	3.7716	3.7716	3.7716		0.0	3.7716	0.00000%
15	79	100	3.8572	3.8574	3.8572		-0.2	3.8572	0.00000%
16	111	100	3.6931	3.6929	3.6931		0.2	3.6931	0.00000%
17	53	100	3.8009	3.8009	3.8009		0.0	3.8009	0.00000%
18	74	100	3.8078	3.8079	3.8079		0.0	3.8079	-0.00263%
19	52	100	3.7902	3.7896	3.7900		0.4	3.7900	0.00528%

Table A1: Environmental Express TDS Stable-Weigh Tests Blank

Analyst	Edward F. Askew								
Date	4/14/2016								
Blank									
Sample #	Bag #	Volume (ml)	Initial weight (g)	Final weight 1 (g)	Final weight 2 (g)	Final weight 3 (g)	Two Consecutive Weights Difference (mg)	Final weight used (mg)	Percent Change
20	80	100	3.7825	3.7826	3.7825		-0.1	3.7825	0.00000%
21	143	100	3.7904	3.7907	3.7904		-0.3	3.7904	0.00000%
22	42	100	3.7561	3.7566	3.7566		0.0	3.7566	-0.01331%
23	141	100	3.8124	3.8124	3.8125		0.1	3.8125	-0.00262%
24	140	100	3.7322	3.7327	3.7324		-0.3	3.7324	-0.00536%
25	139	100	3.7765	3.7763	3.7761		-0.2	3.7761	0.01059%
								Average	-0.00002%
								Standard Deviation	0.00508%

Table A2: Environmental Express TDS Stable-Weigh Tests 200 mg

Analyst	Edward F. Askew									
Date	4/18/2016									
200.03 mg per 50 mL										
Sample #	Bag #	Volume (ml)	Initial weight (g)	Final weight 1 (g)	Final weight 2 (g)	Final weight 3 (g)	Two Consecutive Weights Difference (mg)	Final weight used (mg)	Final Solids Recovered Weight (mg)	Percent Recovery
1	188	50	3.7495	3.9508	3.9505		-0.3	3.9505	201.0	100.50%
2	187	50	3.7712	3.9721	3.9718		-0.3	3.9718	200.6	100.30%
3	186	50	3.8070	4.0078	4.0076		-0.2	4.0076	200.6	100.30%
4	185	50	3.7333	3.9339	3.9338		-0.1	3.9338	200.5	100.25%
5	184	50	3.8077	4.0074	4.0073		-0.1	4.0073	199.6	99.80%
6	182	50	3.7256	3.9263	3.9260		-0.3	3.9260	200.4	100.20%
7	183	50	3.7304	3.9318	3.9314		-0.4	3.9314	201.0	100.50%
8	181	50	3.6925	3.8962	3.8926		-3.6	3.8926	200.1	100.05%
9	180	50	3.7520	3.9522	3.9519		-0.3	3.9519	199.9	99.95%
10	179	50	3.7577	3.9580	3.9585		0.5	3.9585	200.8	100.40%
11	178	50	3.7507	3.9509	3.9506		-0.3	3.9506	199.9	99.95%
12	177	50	3.7350	3.9356	3.9360		0.4	3.9360	201.0	100.50%
13	153	50	3.7772	3.9776	3.9775		-0.1	3.9775	200.3	100.15%
14	165	50	3.8030	4.0027	4.0026		-0.1	4.0026	199.6	99.80%
15	176	50	3.7986	3.9986	3.9985		-0.1	3.9985	199.9	99.95%
16	167	50	3.7678	3.9679	3.9679		0.0	3.9679	200.1	100.05%
17	168	50	3.8320	4.0326	4.0324		-0.2	4.0324	200.4	100.20%
18	166	50	3.7805	3.9820	3.9816		-0.4	3.9816	201.1	100.55%
19	169	50	3.7557	3.9559	3.9557		-0.2	3.9557	200.0	100.00%

Table A2: Environmental Express TDS Stable-Weigh Tests 200 mg

Analyst	Edward F. Askew									
Date	4/18/2016									
200.03 mg per 50 mL										
Sample #	Bag #	Volume (ml)	Initial weight (g)	Final weight 1 (g)	Final weight 2 (g)	Final weight 3 (g)	Two Consecutive Weights Difference (mg)	Final weight used (mg)	Final Solids Recovered Weight (mg)	Percent Recovery
20	171	50	3.7430	3.9435	3.9439		0.4	3.9439	200.9	100.45%
21	170	50	3.8467	4.0469	4.0467		-0.2	4.0467	200.0	100.00%
22	172	50	3.7228	3.9229	3.9228		-0.1	3.9228	200.0	100.00%
23	174	50	3.7589	3.9589	3.9585		-0.4	3.9585	199.6	99.80%
24	138	50	3.8743	4.0761	4.0757		-0.4	4.0757	201.4	100.70%
25	163	50	3.7442	3.9456	3.9458		0.2	3.9458	201.6	100.80%
								Average	200.4	100.21%
								Standard Deviation	0.6	0.29%

Table A3: Environmental Express TDS Stable-Weigh Tests 100 mg

Analyst	Edward F. Askew									
Date	4/20/2016									
100.0 mg per 50 mL										
Sample #	Bag #	Volume (ml)	Initial weight (g)	Final weight 1 (g)	Final weight 2 (g)	Final weight 3 (g)	Two Consecutive Weights Difference (mg)	Final weight used (mg)	Final Solids Recovered Weight (mg)	Percent Recovery
1	160	50	3.8295	3.9296	3.9299		0.3	3.9299	100.4	100.40%
2	159	50	3.8089	3.9089	3.9090		0.1	3.9090	100.1	100.10%
3	157	50	3.8275	3.9276	3.9274		-0.2	3.9274	99.9	99.90%
4	155	50	3.7905	3.8908	3.8912		0.4	3.8912	100.7	100.70%
5	132	50	3.6963	3.7967	3.7964		-0.3	3.7964	100.1	100.10%
6	126	50	3.7543	3.8542	3.8541		-0.1	3.8541	99.8	99.80%
7	121	50	3.8094	3.9093	3.9092		-0.1	3.9092	99.8	99.80%
8	154	50	3.7894	3.8895	3.8898		0.3	3.8898	100.4	100.40%
9	135	50	3.7178	3.8179	3.8177		-0.2	3.8177	99.9	99.90%
10	137	50	3.7489	3.8488	3.8485		-0.3	3.8485	99.6	99.60%
11	136	50	3.7907	3.8907	3.8904		-0.3	3.8904	99.7	99.70%
12	61	50	3.8120	3.9121	3.9120		-0.1	3.9120	100.0	100.00%
13	56	50	3.7894	3.8890	3.8889		-0.1	3.8889	99.5	99.50%
14	NA	50	3.7348	3.8346	3.8344		-0.2	3.8344	99.6	99.60%
15	69	50	3.7634	3.8644	3.8648		0.4	3.8648	101.4	101.40%
16	92	50	3.8360	3.9362	3.9362		0.0	3.9362	100.2	100.20%
17	96	50	3.7731	3.8732	3.8732		0.0	3.8732	100.1	100.10%
18	60	50	3.7506	3.8504	3.8505		0.1	3.8505	99.9	99.90%
19	93	50	3.7791	3.8791	3.8790		-0.1	3.8790	99.9	99.90%

Table A3: Environmental Express TDS Stable-Weigh Tests 100 mg

Analyst	Edward F. Askew									
Date	4/20/2016									
100.0 mg per 50 mL										
Sample #	Bag #	Volume (ml)	Initial weight (g)	Final weight 1 (g)	Final weight 2 (g)	Final weight 3 (g)	Two Consecutive Weights Difference (mg)	Final weight used (mg)	Final Solids Recovered Weight (mg)	Percent Recovery
20	103	50	3.7646	3.8638	3.8635		-0.3	3.8635	98.9	98.90%
21	84	50	3.7944	3.8945	3.8940		-0.5	3.8940	99.6	99.60%
22	78	50	3.8620	3.9630	3.9629		-0.1	3.9629	100.9	100.90%
23	86	50	3.7344	3.8343	3.8341		-0.2	3.8341	99.7	99.70%
24	87	50	3.7374	3.8379	3.8382		0.3	3.8382	100.8	100.80%
25	161	50	3.7346	3.8346	3.8344		-0.2	3.8344	99.8	99.80%
								Average	100.0	100.03%
								Standard Deviation	0.5	0.52%

Table A4: Environmental Express TDS Stable-Weigh Tests 20 mg

Analyst	Edward F. Askew									
Date	4/24/2016									
20.1 mg per 50 mL										
Sample #	Bag #	Volume (ml)	Initial weight (g)	Final weight 1 (g)	Final weight 2 (g)	Final weight 3 (g)	Two Consecutive Weights Difference (mg)	Final weight used (mg)	Final Solids Recovered Weight (mg)	Percent Recovery
1	195	50	3.8442	3.8635	3.8632		-0.3	3.8632	19.0	94.53%
2	194	50	4.1223	4.1430	4.1431		0.1	4.1431	20.8	103.48%
3	193	50	3.9300	3.9498	3.9498		0.0	3.9498	19.8	98.51%
4	191	50	3.8884	3.9087	3.9087		0.0	3.9087	20.3	101.00%
5	189	50	3.9631	3.9836	3.9835		-0.1	3.9835	20.4	101.49%
6	190	50	3.7477	3.7681	3.7680		-0.1	3.7680	20.3	101.00%
7	184	50	3.8975	3.9170	3.9174		0.4	3.9174	19.9	99.00%
8	188	50	3.9284	3.9479	3.9482		0.3	3.9482	19.8	98.51%
9	187	50	3.8906	3.9112	3.9109		-0.3	3.9109	20.3	101.00%
10	186	50	3.9480	3.9680	3.9685		0.5	3.9685	20.5	101.99%
11	209	50	3.8724	3.8925	3.8929		0.4	3.8929	20.5	101.99%
12	185	50	3.9345	3.9543	3.9545		0.2	3.9545	20.0	99.50%
13	210	50	3.9015	3.9221	3.9217		-0.4	3.9217	20.2	100.50%
14	219	50	3.7434	3.7632	3.7635		0.3	3.7635	20.1	100.00%
15	213	50	3.7270	3.7472	3.7473		0.1	3.7473	20.3	101.00%
16	212	50	3.9431	3.9632	3.9633		0.1	3.9633	20.2	100.50%
17	218	50	3.7349	3.7549	3.7551		0.2	3.7551	20.2	100.50%
18	217	50	3.8864	3.9062	3.9064		0.2	3.9064	20.0	99.50%
19	216	50	3.9130	3.9329	3.9331		0.2	3.9331	20.1	100.00%

Table A4: Environmental Express TDS Stable-Weigh Tests 20 mg

Analyst	Edward F. Askew									
Date	4/24/2016									
20.1 mg per 50 mL										
Sample #	Bag #	Volume (ml)	Initial weight (g)	Final weight 1 (g)	Final weight 2 (g)	Final weight 3 (g)	Two Consecutive Weights Difference (mg)	Final weight used (mg)	Final Solids Recovered Weight (mg)	Percent Recovery
20	215	50	3.7322	3.7522	3.7524		0.2	3.7524	20.2	100.50%
21	214	50	3.7237	3.7437	3.7437		0.0	3.7437	20.0	99.50%
22	211	50	3.7281	3.7486	3.7482		-0.4	3.7482	20.1	100.00%
23	220	50	3.7186	3.7394	3.7393		-0.1	3.7393	20.7	102.99%
24	221	50	3.7429	3.7629	3.7629		0.0	3.7629	20.0	99.50%
25	196	50	3.6729	3.6929	3.6931		0.2	3.6931	20.2	100.50%
								Average	20.2	100.28%
								Standard Deviation	0.3	1.72%

Table A5: Environmental Express TDS Stable-Weigh Tests 20 mg-Evaporation Dish

Analyst	Edward F. Askew									
Date	4/22/2016									
20.2 mg per 50 mL (Evaporation Dish)										
Sample #	Evaporation Dish #	Volume (ml)	Initial weight (g)	Final weight 1 (g)	Final weight 2 (g)	Final weight 3 (g)	Two Consecutive Weights Difference (mg)	Final weight used (mg)	Final Solids Recovered Weight (mg)	Percent Recovery
1	1	50	80.0804	80.1007	80.1004		-0.3	80.1004	20.0	99.01%
2	2	50	88.0705	88.0913	88.0916		0.3	88.0916	21.1	104.46%
3	3	50	80.7484	80.7695	80.7698		0.3	80.7698	21.4	105.94%
4	4	50	80.6551	80.6759	80.6758		-0.1	80.6758	20.7	102.48%
5	5	50	80.1832	80.2041	80.2045		0.4	80.2045	21.3	105.45%
6	6	50	71.8200	71.8405	71.8410		0.5	71.8410	21.0	103.96%
7	7	50	77.3487	77.3690	77.3694		0.4	77.3694	20.7	102.48%
8	8	50	71.2989	71.3192	71.3190		-0.2	71.3190	20.1	99.50%
9	9	50	71.0469	71.0674	71.0670		-0.4	71.0670	20.1	99.50%
10	10	50	70.3869	70.4077	70.4072		-0.5	70.4072	20.3	100.50%
11	11	50	71.0414	71.0622	71.0618		-0.4	71.0618	20.4	100.99%
12	12	50	71.3383	71.3591	71.3589		-0.2	71.3589	20.6	101.98%
13	13	50	71.7408	71.7612	71.7612		0.0	71.7612	20.4	100.99%
14	14	50	70.1571	70.1774	70.1776		0.2	70.1776	20.5	101.49%
15	15	50	82.6046	82.6254	82.6256		0.2	82.6256	21.0	103.96%
16	16	50	70.3772	70.3975	70.3976		0.1	70.3976	20.4	100.99%
17	17	50	71.1090	71.1291	71.1294		0.3	71.1294	20.4	100.99%
18	18	50	69.7257	69.7460	69.7464		0.4	69.7464	20.7	102.48%
19	19	50	88.4661	88.4867	88.4872		0.5	88.4872	21.1	104.46%

Table A5: Environmental Express TDS Stable-Weigh Tests 20 mg-Evaporation Dish

Analyst	Edward F. Askew									
Date	4/22/2016									
20.2 mg per 50 mL (Evaporation Dish)										
Sample #	Evaporation Dish #	Volume (ml)	Initial weight (g)	Final weight 1 (g)	Final weight 2 (g)	Final weight 3 (g)	Two Consecutive Weights Difference (mg)	Final weight used (mg)	Final Solids Recovered Weight (mg)	Percent Recovery
20	20	50	91.8973	91.9196	91.9191		-0.5	91.9191	21.8	107.92%
21	21	50	92.1109	92.1311	92.1312		0.1	92.1312	20.3	100.50%
22	22	50	93.7274	93.7480	93.7478		-0.2	93.7478	20.4	100.99%
23	23	50	92.0551	92.0755	92.0750		-0.5	92.0750	19.9	98.51%
24	24	50	94.5897	94.6106	94.6105		-0.1	94.6105	20.8	102.97%
25	A	50	83.3947	83.4134	83.4134		0.0	83.4134	18.7	92.57%
								Average	20.6	101.80%
								Standard Deviation	0.6	2.99%

Table A6: StableWeigh and Porcelain Evaporation Dish Precision

TDS Mass	Vessel	Average	Standard Deviation	% RSD	Duplicate Relative Percent Difference
200 mg	StableWeigh	200.4	0.57	0.29%	1.00%
100 mg	StableWeigh	100.0	0.52	0.52%	2.50%
20 mg	StableWeigh	20.2	0.35	1.71%	9.05%
20 mg	Porcelain Evaporation Dish	20.6	0.60	2.94%	15.31%

Table A7: StableWeigh Method Blank

Sample #	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25
Blank Weight Difference (mg)	-0.1	0	-0.1	0.2	-0.2	0	-0.1	0	0	-0.3	0.4	0	-0.1	0	0	0	0	0.1	-0.2	0	0	0.5	0.1	0.2	-0.4

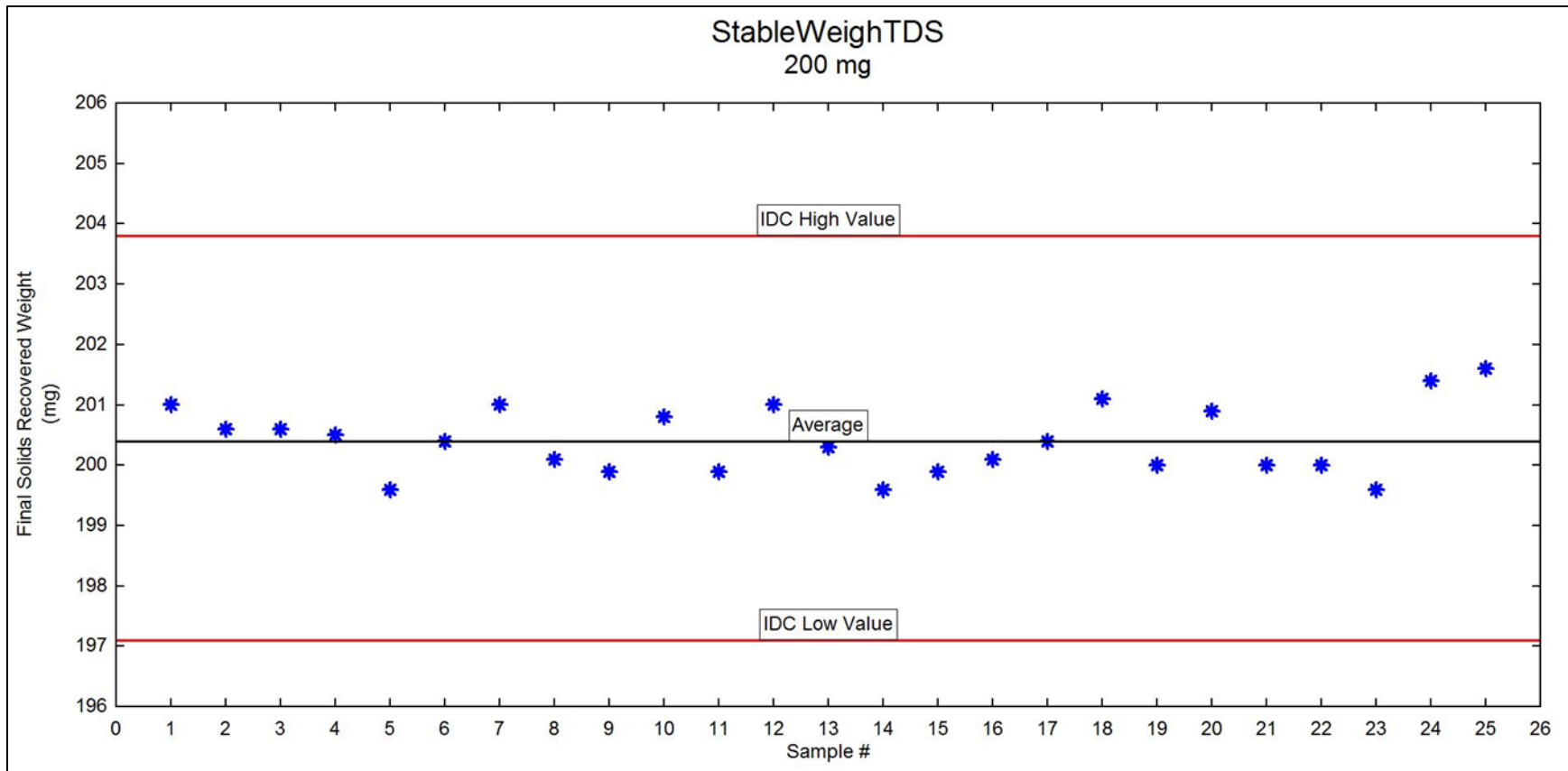


Figure 6: StableWeigh 200 mg Initial Demonstration of Capability

Stableweigh TDS 100 mg

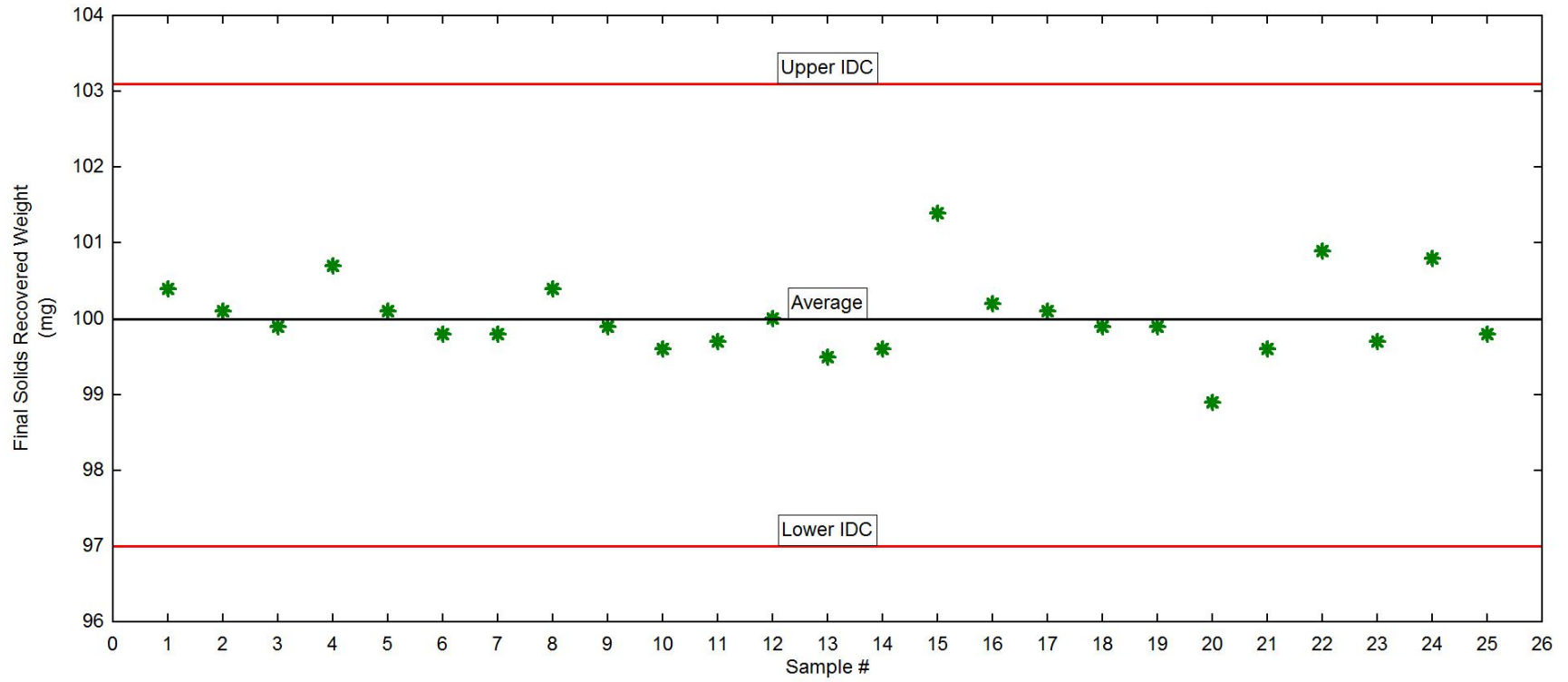


Figure 7: StableWeigh 100 mg Initial Demonstration of Capability

TDS
20 mg

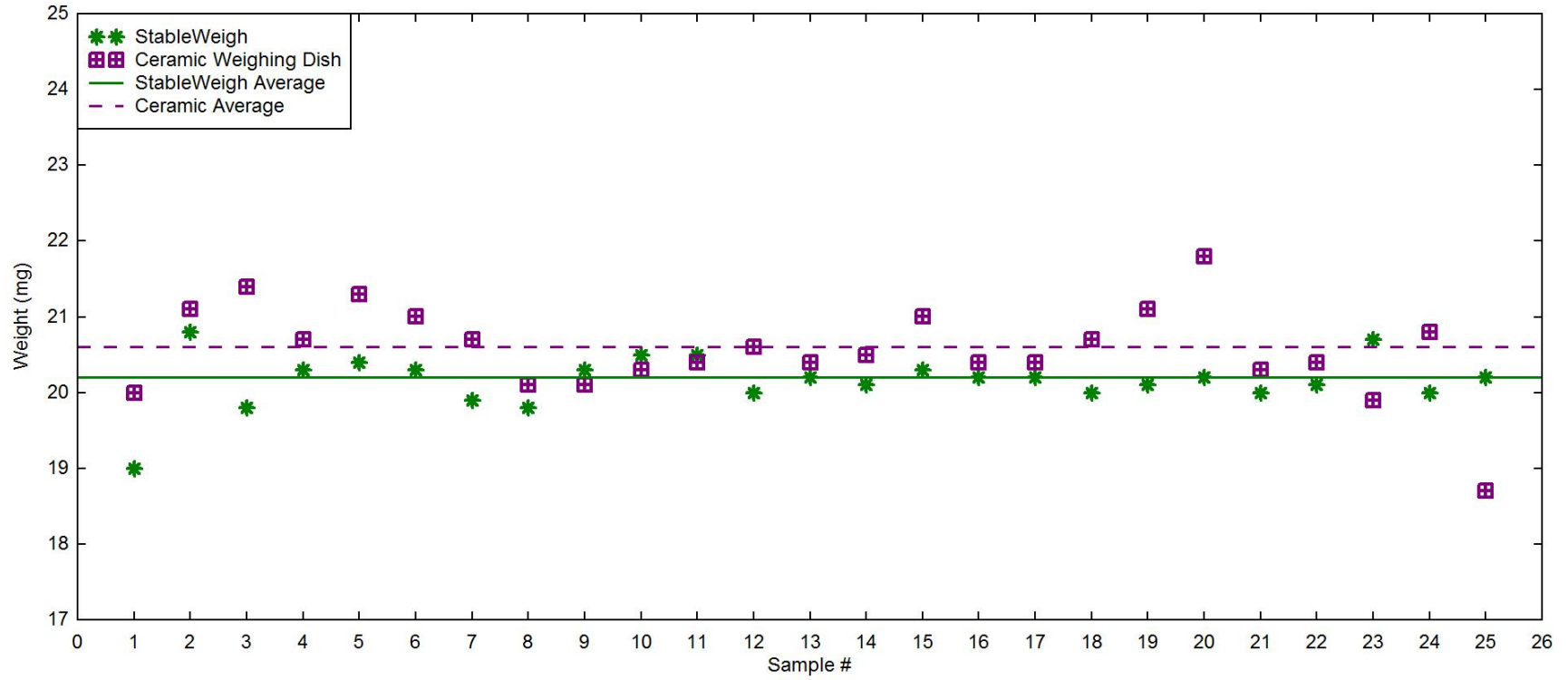


Figure 8: StableWeigh vs. Porcelain Evaporation Dish Averages