

	Environmental Analysis Teaching and Research Laboratory	Date: 8/11/2016	Number: 23
	Standard Operating Procedure	Title: SOP for In Vitro Determination of Chlorophyll <i>a</i> Concentrations by Fluorescence	
	Approved By: Los Huertos	Revision Date: / /	

1. Scope and Application

1.1 This method provides a procedure for the fluorometric determination of chlorophyll *a* and its magnesium-free derivative, pheophytin *a* in marine and freshwater phytoplankton.

1.2 This method is modified from the US EPA Method 445.0 and APHA Standard Methods for the Examination of Water and Wastewater, 22nd Edition.

2. Summary of Method

2.1 Chlorophyll-containing phytoplankton in a measured volume of sample water are concentrated by filtering at low vacuum (13 cm Hg) through a glass fiber filter (Whatman GF/F). The pigments are extracted from the phytoplankton in 90% acetone and to ensure thorough extraction of chlorophyll *a*, are allowed to steep for at least 2hrs. The fluorescence of the sample is measured at the excitation wavelength of 485 nm and emission wavelengths 685 / 50 nm. Sample fluorescence is measured before and after acidification with 0.1M HCl to obtain a corrected chlorophyll *a* concentration.

3. Definitions

3.1 Stock Standard Solution (SSS) – a solution prepared in the laboratory using reference materials purchased from a reputable commercial source.

3.2 Laboratory Reagent Blank (LRB) – an aliquot of reagent water (Milli-Q/SuperQ) or other blank matrices that are treated the same as the sample including exposure to all glassware, equipment, solvents, reagents, internal standards and surrogates that are used with other samples. The LRB is used to determine if method analytes or other interferences are present in the laboratory environment, reagents or apparatus.

3.3 Field duplicates– Two separate samples collected at the same time and placed under identical circumstances and treated exactly the same throughout the field and laboratory procedures. Provide a measure of the precision associated with sample collection, preservation, storage and laboratory processing.

3.4 Quality Control Sample (QCs)– A solution of known concentration obtained from a source external to the laboratory to check laboratory performance.

4. Interferences

4.1 Any substance extracted from the filter or acquired from laboratory contamination that fluoresces in the red region of the spectrum may interfere in the accurate measurement of both chlorophyll *a* and pheophytin *a*.

4.2 item Spectral interferences resulting from the fluorescence of the accessory pigment chlorophyll *b*, and the chlorophyll *a* degradation product pheophytin *a*, can result in the overestimation of chlorophyll *a* concentrations. However, highly selective optical filters used in this method minimize these interferences.

4.3 Quenching effects are observed in highly concentrated solutions or in the presence of high concentrations of other chlorophylls and carotenoids. Samples should be diluted.

4.4 Fluorescence is temperature dependent with higher sensitivity occurring at lower temperatures. Samples, standards, LRBS (section 3.2) and QCs (section 3.4) must be at the same temperature to prevent errors and maximize precision. Analysis of samples at ambient temperatures is required in this method.

4.5 All photosynthetic pigments are light and temperature sensitive. Work must be performed in subdued light and all standards, QC materials and filter samples must be stored in the dark at -20°C to prevent degradation.

5. Safety and Personnel Protective Equipment

5.1 Lab safety-glasses and lab coats are required for all laboratory analysis. Use gloves to avoid skin irritation from contact with acetone; work under fume hood when possible.

5.2 Sonication Procedure – When using the sonicator always wear ear protection. **Sonicator should never be used outside of solution.**

5.3 Chemical Hygiene – Please refer to the Material Data Safety Sheets (MSDS) files for questions concerning a chemical's toxicity and the necessary safety precautions.

5.4 Waste Disposal– Dispose of waste in the acetone collection bottle. See "SOP 02 Handling of Hazardous Materials."

6. Related Documents

Other SOPs that are currently be developed!

- "SOP 02 for Handling of Hazardous Materials"
- "SOP 10 for using Balances, Pipettes and Glassware"
- "SOP 11 for preparing Standards and QCs"

7. Materials and Apparatus

7.1 Turner Designs P/N 998-7210 Fluorometer

7.2 10mL borosilicate glass tubes with caps (**Cat #?**)

7.3 Whatmann glass micro fiber filter GF/F 0.7 μ m retention 47mm (**Cat #?**)

7.4 Millipore glass filtration unit with vacuum and 47mm fritted glass disk base (**Cat #?**)

7.5 Tweezers or flat tipped forceps (**Cat #?**)

7.6 Assorted class A calibrated pipettes (**Cat #?**)

7.7 50mL, 100mL, and 1-L class A volumetric flask

7.8 Sonicator

7.9 Centrifuge (specs)

7.10 15 mL centrifuge tubes? (**Cat #?**)

7.11 Note(for lab techs): All reusable labware that comes in contact with chlorophyll solutions should be cleaned with acetone and acid free. Diswashing should include soaking in laboratory grade detergent and water, rinsing with tap water then rinsing with deionized water.

8. Reagents and Standards

8.1 Acetone, HPLC grade

8.2 Hydrochloric Acid

8.3 Chlorophyll *a* free of chlorophyll *b*. May be obtained from a commercial supplier.

8.4 Milli-Q/Super-Q water

8.5 **0.1 M HCl solution**– Add 0.85 mL of concentrated HCl to approximately 50mL of water and dilute to 100 mL.

8.6 **Aqueous Acetone Solution**– 90% Acetone/ 10% Milli-Q water. Carefully measure 100mL of water into a 100mL volumetric flask. Then pour the water into a 1L volumetric flask. Fill the 1L flask up to the line with acetone. Mix and then transfer to a dark amber bottle. Label the bottle and store in the flammables cabinet (under the middle fumehood in the Los Huertos Lab).

8.7 **Chlorophyll Standard Stock Solution (SSS)**- Chlorophyll *a* from Sigma is shipped in an amber glass ampoule. This should be stored in the freezer until use. Tap the ampoule

until all of the dried chlorophyll has settled on the bottom. Working in a darkened room, carefully break the tip off the ampoule and transfer the contents into a 50mL volumetric flask and dilute to volume with 90% acetone (from section 8.6). Transfer to a darkened bottle or wrap the flask in foil to protect it from light. The concentration of the solution must be determined spectrophotometrically using a multiwavelength spectrophotometer. Label bottle including the chlorophyll lot number. When stored in an airtight container at room temperature, the SSS is stable for at least six months.

8.8 Chlorophyll *a* Primary Dilution Standard (PDS)- Add 1mL of the SSS (section 8.7) to a 100mL volumetric flask and dilute to volume with aqueous acetone solution (section 8.6). If exactly 1mg of pure chlorophyll was used to prepare the SSS, the concentration of the PDS is 200 µg/L. Prepare fresh prior to use and label flask and wrap in foil.

8.9 Quality Control Samples (QCs)- Since there are no commercially available QCs, they must be prepared from the PDS at the following concentrations:

- QC1 = 5 µg/L
- QC1 = 20 µg/L
- QC1 = 50µg/L
- QC1 = 200 µg/L

9. Extraction

9.1 Invert the sample bottle gently several times before filtering to suspend particles. Mixing the sample well improves the replication greatly.

9.2 Conduct filtration in an area with subdued light. Gently filter sample, filter vacuum should not exceed 20 kPa). Sample volume to be filtered will vary depending on the algal concentration in the sample. The sample needs to be filtered until a green tint can be seen on the filter, filter sample in 20mL increments using a graduated cylinder or calibrated pipette. For benthic samples a much smaller volume will need to be filtered, roughly 5mL, freshwater sample needed to be filtered can range from 20mL to 300mL, and seawater sample volume can exceed 1 L. It is extremely important to record the filtered sample volume during this process. If the filter is too green the sample analysis will be over range and will need to be diluted, however it is best to avoid diluting the sample when possible.

9.3 After filtration, fold each filter in half, using tweezers and a spatula and a place inside a clean 50mL falcon tube. Perform a laboratory duplicate and blank (100mL Milli-Q) every 10-15 samples.

9.4 Freeze filter samples for up to 28 days before extraction

9.5 To extract chlorophyll *a*, add 25 mL of 90% aqueous acetone solution to the falcon tubes ensuring each filter is fully submerged in the solvent. Cap the tubes and invert several times

to mix. Turn on and set sonicator dial to 7. Sonicate by sumberg the tip of the sonicator about halfway into the acetone solution and pressing the trigger. To help break up the filter completely, be sure to move tip around the entire sample. After sonication, vigorously shake sample several times, cover with foil and then steep in freezer for at least 2 hrs.

9.6 Remove samples from the freezer and allow both the sample and QCs to come to room temperature.

9.7 Mash sample filters to release any chlorophyll that has accumulated on the filter. Remove and dispose of the filter using forceps.

9.8 Pour the filter extracts into 15mL falcon tubes and label. Then centrifuge them at (insert the right settings for centrifuge here) for 3 minutes. Or until all filter material is separated from the extration liquid. When removing the tubes from the centrifuge, be careful not to re-suspend filter particles from the bottom of the tubes.

10. Sample Analysis

10.1 2.5 mL of each sample will be dispensed into a clean glass cuvette, again taking care not to resuspend filter particles.

10.2 Turn on the fluorometer (back left side) and allow 10 minutes to for warmup countdown to complete. Turn on lab computer and open SIS for Turner software to record data.

10.3 Use "Measure Raw Fluorescence" to take sample measurements.

10.4 Use the 90% acetone solution for a blank measurement on the sensitivity setting that will be used for sample analysis.

10.5 Read and record the values for the QCs prior to sample analysis, the Turner Fluorometer will send and accumulate data to the connected computer using the "SIS for Trilogy" program.

10.6 Wipe the outside of the cuvette with a kimwipe before taking a measurement then measure the raw fluorecence of the sample. If the chlorophyll *a* concentration of the sample is 90% of the upper limit of the LDR (LDR=500 µg/L) it will be necessary to perform a dilution. However, samples should not be over diluted so be sure to use an appropriate dilution factor.

For a 1:2 Dilution This dilution will be the one most likely needed in the case above. Simply add a 1 mL aliquot of the sample into 1 mL aqueous acetone solution.

For a 1:100 Dilution If the concentration of chlorophyll *a* in the sample is very high you can try this dilution factor. Add 50 µL of sample to 4.95 mL of aqueous acetone solution.

10.7 It will be necessary to calculate corrected chlorophyll *a* values to determine wether

dilutions need to be made. This is because pheophytin *a* concentrations will not be the same across all samples. An excel spreadsheet template with equations can be found on

10.8 After reading the fluorescence of all samples, add 75 µL of 0.1M HCl to each tube. Thoroughly mix the sample by sealing cuvette with a rubber stopper and inverting several times. Allow 90 seconds before measuring the fluorescence of all the samples again.

11. Data Analysis and Calculations

To calculate 'corrected chlorophyll *a*' values use the following equations:

11.1 Variables

1. $[Std]$ = Expected Standard concentration (This is not calculated, use the concentration of one of the standards that are run with the samples and its corresponding F value)
2. S_b = Standard reading before acidification
3. S_a Standard reading after acidification
4. F_b reading before acidification
5. F_a reading after acidification
6. V_e volume of extract
7. V_s volume of sample

11.2 Equations

Equation 1: Acid Ratio (AR) = $(S_b - \text{Blank}) / (S_a - \text{Blank})$

Equation 2: Interpolation Before (InterpB) = $[Std] * (F_b - \text{Blank}) / (S_b - \text{Blank})$

Equation 3: Interpolation After (InterpA) = $[Std] * (F_a - \text{Blank}) / (S_b - \text{Blank})$

Equation 4: Corrected Chlorophyll *a* = $[AR / (AR - 1)] * (\text{InterpB} - \text{InterpA}) * (V_e / V_s)$

Equation 5: Corrected Pheophytin *a* = $((AR * \text{InterpA}) - \text{InterpB}) * (V_e / V_s)$

Note: Generally, equation 5 is not used.

12. QC/QA Criteria

12.1 An LRB is performed every 10 samples. The LRB should be less than the calculated lower detection limit for analysis.

12.2 The QCs should be within $\pm 20\%$ of the known value.

12.3 The RSD of the replicate measurements should be within $\pm 20\%$