

# Sediment Organic Matter

v 0.1.3



<u>How to cite this work:</u> Seagrass Habitats: MarineGEO Protocols. (2019) Tennenbaum Marine Observatories Network, MarineGEO, Smithsonian Institution. DOI:





# Introduction

This protocol provides standardized data on sediment bulk density and organic matter content, which is obtained through loss-on-ignition using a combustion furnace.

Additional copies of this protocol, field datasheets, data entry templates, instructional videos, literature, and more can be found on the MarineGEO protocol website: <a href="https://marinegeo.github.io/">https://marinegeo.github.io/</a>.

#### **Measured Parameters**

This protocol quantifies the organic matter content in marine sediments, measured as:

- Bulk density (g/mL) \*if possible
- Sediment dry mass (g)
- Sediment ash-free dry mass (g)

# Requirements\*

\*Estimated times will vary by site and conditions

Personnel: 2 people

Estimated Total Time Per Site (i.e., all three locations at the site):

Preparation: 1 person x 1 day Field work: 2 people x 1 day

Post processing: 1 person x 3-5 days Data processing: 1 person x 1 day

Replication: Three (3) 5 cm-x-5 cm sediment cores taken along three (3) transects (total n = 9)

Materials:

Sui	rvey Design:
	1 50-m metric transect tape
	Hand-held GPS unit
	2 PVC marker poles (diameter and length as needed)
<u>Fie</u>	<u>ldwork:</u>
	9 small plastic bags with external and internal labels



9 5-mL plastic syringes with graduations (0.1-0.2 mL) with the applicator tip cut off 1 cooler with ice (optional)
st-processing: 9 pre-weighed foil tins
Pencil/pen
Drying oven
Combustion furnace

#### Methods

Fully review this and any additional protocols necessary for the sampling excursion. Address any questions or concerns to <a href="mailto:marinegeo@si.edu">marinegeo@si.edu</a> before beginning this protocol.

### Preparation:

- 1. Review the MarineGEO <u>Seagrass Habitats Survey Design</u> for site selection and setup. This protocol assumes n = 3 sediment samples taken every 10-12 m along a 50-m transect, replicated along 3 separate transects.
- 2. Label 9 disposable plastic bags with the sampling location, transect, and replicate number using a permanent marker.
- 3. Place a plastic syringe and an internal label with the same metadata written on waterproof paper inside the corresponding plastic bag.
- 4. Fill a cooler with ice immediately before departing for the field.

#### Fieldwork:

- 1. At each replicate along the transect, randomly select an unvegetated patch ~1 m to any side of the transect.
- 2. Remove the plunger from the syringe. Take the open end of the 50-mL syringe and gently insert it into the sediment to a depth of ~5-10 cm. Take care to avoid any structures like rhizomes or woody debris.
- 3. Place the plunger into the syringe to create suction, and then gently extract the syringe from the sediment.
- 4. Place the syringe with the trapped sediment into the plastic bag and seal it.
- 5. Repeat steps 1-4 at the next location along the first transect until all 3 replicates are taken.
- 6. Repeat steps 1-5 for the remaining two transects for a total of 9 samples.
- 7. Place all bags on ice in the cooler. Transport cooler with samples back to the lab for processing.

#### Post-processing:



Samples are best processed immediately (within 24 hours) of returning from the field. Samples can be stored for longer in the refrigerator, but risks evaporation.

- 1. Print lab data sheets.
- 2. Weigh foil tins and record the weight of the tin directly on the foil using a pen. Tins can be either pre-made or constructed by folding an aluminum foil square over on itself and sealing the sides.
- 3. Select a replicate syringe and push the plunger to discard all but the top 5 cm of sediment.
- 4. If the samples did not take on or dissolve in water and did not dry out, record the volume of sediment (to the nearest 0.1 mL).
- 5. Use the plunger to push the sediment plug into a pre-weighed tin.
- 6. Remove all visible fauna, large shells, rhizomes/roots/woody debris from the sample. Work quickly to minimize loss of water.
- 7. Place the labeled foil tins in a drying oven at 60°C and dry to constant weight (usually 1-3 days, depending on the volume of material).
- 8. Weigh the tin and dried sediment plug, and record the dry weight on the lab data sheet
- 9. Combust the samples at 520°C for 5 hours.
- 10. Let the sample cool in the drying oven to avoid taking on any moisture, then weigh the tin and combusted sediment plug, and record the ash-free dry weight on the lab data sheet.

# **Data Submission**

- 1. Enter data into provided data entry template.
- 2. Scan the completed lab data sheets and save both paper and electronic versions.
- 3. E-mail data entry file and scanned lab data sheets to: <a href="mailto:marinegeo-data@si.edu">marinegeo-data@si.edu</a>.