

# Transparent Exopolymer Particles (TEP) from CTD samples collected during R/V Hugh R. Sharp cruise HRS2204 from Apr to May 2022

**Website:** <https://www.bco-dmo.org/dataset/945964>

**Data Type:** Cruise Results

**Version:** 1

**Version Date:** 2024-12-13

## Project

» [Collaborative Research: The importance of particle disaggregation on biogeochemical flux predictions](#)

(Disaggregation)

Contributors	Affiliation	Role
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## Abstract

These data include measurements of transparent exopolymer particles (TEP) from CTD bottle water samples collected during a cruise on the Northeast Continental Shelf to study particle disaggregation. One cruise was completed aboard the R/V Hugh R. Sharp from 2022-04-21 through 2022-05-02 (HRS22-04), which visited a variety of stations and hydrodynamic environments associated with the Northeast Continental Shelf of the United States. Stations ranged from Georges Bank and the Great South Channel near the Gulf of Maine, Martha's Vineyard, the mouth of the Sakonnet River near Newport, Rhode Island, and Hudson Canyon near New York. TEP measurements were performed following recently published methods relying on Alcian Blue staining protocols and spectrophotometry. These data were collected as part of a study to clarify the importance of hydrodynamic forces on the cohesion, aggregation, and breakup of marine particles. These data were collected by Dr. Kieran Curran of the University of New Hampshire on the cruise led by Dr. Matthew Rau (chief-sci) of the George Washington University.

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## Coverage

**Location:** Northeastern U.S. Continental Shelf

**Spatial Extent:** N:41.7749 E:-67.3996 S:39.4554 W:-72.2765

**Temporal Extent:** 2022-04-23 - 2022-05-01

## Methods & Sampling

This cruise visited eight stations on the Northeastern U.S. Continental Shelf. Latitudes and longitudes provided per sample in the data, but general station descriptions are below.

- Station 1 - Station 3: Georges Bank near the Gulf of Maine. Approximate location was 41.7 N, 68 W. Samples acquired from 0 m to 25 m depths.
- Station 4: The Great South Channel near the Gulf of Maine. Approximate location was 41.6 N, 69 W. Samples acquired from 0 m to 150 m depths.
- Station 5: Only one CTD profile was taken before this station was aborted due to weather. No data acquired. Station location was 40.8 N, 70.5 W.
- Station 6: Off the coast of Martha's Vineyard. Approximate location was 41.3 N, 70.5 W. Samples acquired from 0 m to 10 m depths.
- Station 7: At the mouth of the Sakonnet River near Newport, Rhode Island. Approximate location was 41.5 N, 71.2 W. Samples acquired from 0 m to 10 m depths.
- Station 8: Hudson Canyon near New York. Approximate location was 39.5 N, 72.3 W. Samples acquired from 0 m to 200 m depths.

Seawater samples between 500 and 2000 ml were vacuum filtered through a 0.45 micron polycarbonate membrane filter. Filter retentate was then stained with Alcian Blue solution. The Alcian Blue solution was made to a concentration of 400 mg/L by mixing dye powder with ultrapure water that was acidified to pH 2.5 with glacial acetic acid. After mixing, the stain was filtered through a 0.22 micron polycarbonate membrane to remove undissolved powder. Stained filters were frozen in Falcon tubes at -20 C for the remainder of the cruise and returned to the lab on dry ice for analysis. In the lab, TEP samples were extracted in 6ml 80% sulfuric acid solution for 2 hrs. The absorbance of this solution was then measured with a spectrophotometer at 787 nm. Concentration of TEP was converted from absorbance to microgram equivalent of Xanthan Gum through a calibration curve. Calibration was conducted following the procedure of Bittar, et al. (2018). A 5-point calibration was performed using dilutions of Xanthan Gum solution with concentrations of 9.37, 18.75, 37.50, 56.25, and 75 micrograms per ml in 1 mL polypropylene tubes. 0.5 ml of the Alcian Blue solution was added to these tubes, which were then mixed for 1 min through manual agitation. These stained aliquots were then poured onto filtration towers and filtered through 0.45 micron polycarbonate filters. The filter retentate was then extracted using the 80% sulfuric acid wash for spectrophotometric quantification. Filters were soaked in the solution for 2-20 hrs with regular agitation. The spectrophotometer (Agilent 8453) was first blanked with ultrapure water prior to measurements. 6 mL of extraction solution was read using 1.0 cm polystyrene disposable cuvettes (Fisher-Scientific).

## Data Processing Description

Microsoft excel was used to create the linear calibration curve from the Xanthan Gum dilutions and to convert measured absorbance into mass of TEP in Xanthan Gum equivalents.

## BCO-DMO Processing Description

- Imported "HRS22-04\_TEP.xlsx" into the BCO-DMO system
- Corrected the import formatting which seems to have left off the seconds for time value on row 39; value is "11:45:10"
- Combined date and times to create datetime field
- Renamed fields to comply with BCO-DMO naming conventions
- Exported final file as "945964\_v1\_tep\_ctd.csv"

## Problem Description

No problems were encountered though TEP samples were not acquired on every depth and CTD cast.

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## Related Publications

Bittar, T. B., Passow, U., Hamaraty, L., Bidle, K. D., & Harvey, E. L. (2018). An updated method for the calibration of transparent exopolymer particle measurements. *Limnology and Oceanography: Methods*, 16(10), 621-628. doi:[10.1002/lom3.10268](https://doi.org/10.1002/lom3.10268)

## Parameters

Parameter	Description	Units
ID	Identifier of the CTD cast sample, formatted as (station#_CTDcast#_depth)	unitless
Station	Station number	unitless
CTD	CTD cast number	unitless
Depth	Depth in meters that the CTD bottle was triggered for sampling	meters (m)
ISO_DateTime_UTC	ISO datetime the sample was acquired in UTC	unitless
Date_UTC	UTC date on which the sample was acquired	unitless
Time_UTC	UTC time when the sample was acquired	unitless
Latitude	Ship's latitude when the sample was taken	decimal degrees
Longitude	Ship's longitude when the sample was taken	decimal degrees
TEP_xg	Mass concentration of TEP in Xanthan Gum equivalent, measured in micrograms per liter. Minimum detection limits were 5.6 micrograms per liter.	micrograms per liter (ug/L)

## Instruments

<b>Dataset-specific Instrument Name</b>	CTD
<b>Generic Instrument Name</b>	CTD Sea-Bird 911
<b>Dataset-specific Description</b>	These data include measurements of transparent exopolymer particles (TEP) from CTD bottle water samples collected during a cruise on the Northeast Continental Shelf to study particle disaggregation.
<b>Generic Instrument Description</b>	The Sea-Bird SBE 911 is a type of CTD instrument package. The SBE 911 includes the SBE 9 Underwater Unit and the SBE 11 Deck Unit (for real-time readout using conductive wire) for deployment from a vessel. The combination of the SBE 9 and SBE 11 is called a SBE 911. The SBE 9 uses Sea-Bird's standard modular temperature and conductivity sensors (SBE 3 and SBE 4). The SBE 9 CTD can be configured with auxiliary sensors to measure other parameters including dissolved oxygen, pH, turbidity, fluorescence, light (PAR), light transmission, etc.). More information from Sea-Bird Electronics.

<b>Dataset-specific Instrument Name</b>	Agilent 8453 UV-visible
<b>Generic Instrument Name</b>	Spectrophotometer
<b>Dataset-specific Description</b>	A spectrophotometer (Agilent 8453 UV-visible) was used to make the absorbance measurements, which were read using 1.0 cm polystyrene disposable cuvettes (Fisher-Scientific).
<b>Generic Instrument Description</b>	An instrument used to measure the relative absorption of electromagnetic radiation of different wavelengths in the near infra-red, visible and ultraviolet wavebands by samples.

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## Deployments

### HRS2204

<b>Website</b>	<a href="https://www.bco-dmo.org/deployment/946038">https://www.bco-dmo.org/deployment/946038</a>
<b>Platform</b>	R/V Hugh R. Sharp
<b>Start Date</b>	2022-04-21
<b>End Date</b>	2022-05-02
<b>Description</b>	See additional cruise information in R2R: <a href="https://www.rvdata.us/search/cruise/HRS2204">https://www.rvdata.us/search/cruise/HRS2204</a>

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## Project Information

**Collaborative Research: The importance of particle disaggregation on biogeochemical flux predictions (Disaggregation)**

**Coverage:** Northeast United States Continental Shelf

NSF abstract:

Particle settling is one of the major ways that material in surface waters reaches the deep ocean. Particulate matter in the open ocean consists primarily of organic material from plankton and other biological detritus, which can readily aggregate to form large flocs. A combination of physical, chemical, and biological processes transforms these flocs as they settle, redistributing material throughout the water column and potentially sequestering elements such as carbon in the deep ocean. The impact of these transformations is affected by the sinking speed of these flocs, with larger and denser particles settling faster than smaller, less-dense ones. One of the key questions facing oceanographers today is what controls particle settling speed (for example, particle size, shape, and density). There is considerable evidence that particles readily break apart as they settle, decreasing their average size and settling speed, but it is not yet understood what conditions cause these disaggregation events. This work will measure the breakup characteristics of organic settling particles both in the laboratory and at sea to quantify the importance of these breakup processes relative to particle transport. The work will be done at the Pennsylvania State University in collaboration with the University of Georgia to target the development of future marine particle disaggregation models for use by the oceanographic community.

This research will play an important role in determining the importance of disaggregation on the vertical transport of particulate matter in the ocean. The project will quantify the breakup of organic marine aggregates due to fluid forces caused by turbulence or swimming organisms. Phytoplankton will be cultured and formed into aggregates in the lab prior to disaggregation using calibrated turbulence. The size, shape, and structure of these aggregates before and after breakup will be quantified using high-speed visualization and holographic imaging. In addition to the laboratory measurements, a deployable instrument that can disrupt particles in-situ and measure their size and shape will be built and deployed in the North Atlantic during the spring bloom of phytoplankton. Detailed measurements of particle concentrations, breakup characteristics, organic content, and ambient turbulence as a function of depth in the water column will be collected. This work will represent the first study of marine aggregate breakup in-situ. Specifically, the project will clarify: (1) under what conditions disaggregation is important, (2) how strong different types of natural marine aggregates are and how their strength varies with size, composition, and morphology, and (3) how aggregate size, composition, and structure influences the distribution of its breakup mass. This project will advance the career of a doctoral student and engage numerous undergraduate researchers with the field of ocean science.

This award reflects NSF's statutory mission and has been deemed worthy of support through evaluation using the Foundation's intellectual merit and broader impacts review criteria.

### **Additional Project Output (supplement to Data Collections section below):**

#### Model Code Description:

Adrian Burd's Research Lab. (2023). BurdLab/Dissaggregation: Disaggregation (Disaggregation). Zenodo. <https://doi.org/10.5281/zenodo.8226166>

Associated Github Repository: <https://github.com/BurdLab/Dissaggregation/tree/Disaggregation>

This is the initial release of model code for particle aggregation and disaggregation in the ocean. The referenced Github Repository contains Matlab code to calculate the evolution of the particle size distribution in a single layer of the water column. The code numerically solves the aggregation-disaggregation mass balance equations using a so-called sectional approach developed by Gelbard and Seinfeld (J. Colloid and Interface Sci., 68:363-382, 1979). The model allows for particle aggregation, disaggregation, and sinking, and also changes in aggregate size from cell growth (see SetupCoag.m), and will form the basis of a suite of particle aggregation/disaggregation models. All documentation is provided within the code itself. Please see Associated Github Repository link above for detailed description and files.

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## **Funding**

<b>Funding Source</b>	<b>Award</b>
<a href="#">NSF Division of Ocean Sciences (NSF OCE)</a>	<a href="#">OCE-1948283</a>

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