

# Photosynthetic data collected from the R/V Oceanus OC1504A in the Oregon/California Coastal Upwelling Zone, between 34-44N and 120-124W in 2015.

**Website:** <https://www.bco-dmo.org/dataset/652739>

**Data Type:** Cruise Results

**Version:** 1

**Version Date:** 2016-07-28

## Project

» [Linking physiological and molecular aspects of diatom silicification in field populations](#) (Diatom Silicification)

Contributors	Affiliation	Role
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<a href="#">Brzezinski, Mark A.</a>	University of California-Santa Barbara (UCSB-LifeSci)	Co-Principal Investigator
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## Abstract

Photosynthetic data collected from the R/V Oceanus OC1504A in the Oregon/California Coastal Upwelling Zone, between 34-44N and 120-124W in 2015.

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## Coverage

**Temporal Extent:** 2015-04-20 - 2015-05-01

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## Dataset Description

Photosynthetic data on water collected by CTD and measured using fast repetition rate fluorometry.

## Methods & Sampling

Photosynthetic parameters were measured using fast repetition rate fluorometry on whole seawater collected by CTD. See reference below for details on data analysis.

## Data Processing Description

Photosynthetic parameters were corrected for background fluorescence by measuring 0.2  $\mu\text{m}$  filtered seawater from 1-2 depths.  $F_o$  and  $F_m$  of background samples were subtracted from sample  $F_o$  and  $F_m$  and corrected values were used to calculate  $F_v/F_m$ , where  $F_v = F_m - F_o$  (Kolber et al. 1998)

**DMO Notes:**

- File was resubmitted by PI after some consultation with several columns and rows removed
- Column names were changed to meet BCO-DMO standards
- Some spaces were removed from cell contents
- cruise\_id and ISO\_DateTime\_UTC column were added

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## Data Files

File
<b>photosynthetic_data.csv</b> (Comma Separated Values (.csv), 4.45 KB) MD5:3a1d21a92492aafc6354c6ec2dcc10c8
Primary data file for dataset ID 652739

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## Related Publications

Kolber, Z. S., Prášil, O., & Falkowski, P. G. (1998). Measurements of variable chlorophyll fluorescence using fast repetition rate techniques: defining methodology and experimental protocols. *Biochimica et Biophysica Acta (BBA) - Bioenergetics*, 1367(1-3), 88-106. doi:[10.1016/s0005-2728\(98\)00135-2](https://doi.org/10.1016/s0005-2728(98)00135-2)  
*General*

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## Parameters

Parameter	Description	Units
cruise_id	cruise identification where samples were collected	unitless
CTD	CTD cast	unitless
depth	depth at which samples were collected	meters
date_local	local date of sample collection; mm/dd/yy	unitless
time_local	local time of sample collection; HH:MM:SSpp	unitless
Fluor_min	minimal fluorescence yield corrected for background fluorescence. $F_o$	relative units
Fluor_max	maximal fluorescence yield corrected for background fluorescence. $F_m$	relative units
FvFm	maximum quantum yield corrected for background fluorescence; $F_v$ divided by $F_m$	dimensionless
functional_absorption	Functional absorption cross-section of photosystem II (measured using 450 nm excitation; units $A_2$ ); sigma	unitless
connectivity_p	connectivity factor defines the efficiency of exciton energy transfer between individual photosynthetic units; originally p	unitless
ISO_DateTime_UTC	DateTime (UTC) ISO formatted	unitless

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## Instruments

<b>Dataset-specific Instrument Name</b>	Fast Repetition Rate Fluorometer
<b>Generic Instrument Name</b>	Fast Repetition Rate Fluorometer
<b>Dataset-specific Description</b>	Photosynthetic parameters were measured.
<b>Generic Instrument Description</b>	An FRRf is used for measuring the fluorescence of a sample of phytoplankton photosynthetic competency (Fv/Fm).

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## Deployments

### OC1504A

<b>Website</b>	<a href="https://www.bco-dmo.org/deployment/560135">https://www.bco-dmo.org/deployment/560135</a>
<b>Platform</b>	R/V Oceanus
<b>Report</b>	<a href="https://musicc2015.wordpress.com">https://musicc2015.wordpress.com</a>
<b>Start Date</b>	2015-04-19
<b>End Date</b>	2015-05-02
<b>Description</b>	Data for the project "Linking physiological and molecular aspects of diatom silicification in field populations" (PIs Kimberlee Thamatrakoln and Mark Brzezinski) were collected on this cruise.

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## Project Information

### Linking physiological and molecular aspects of diatom silicification in field populations (Diatom Silicification)

**Coverage:** Oregon/California Coastal Upwelling Zone, between 34-44N and 120-124W

#### *Description from NSF award abstract:*

Diatoms, unicellular, eukaryotic photoautotrophs, are among the most ecologically successful and functionally diverse organisms in the ocean. In addition to contributing one-fifth of total global primary productivity, diatoms are also the largest group of silicifying organisms in the ocean. Thus, diatoms form a critical link between the carbon and silicon (Si) cycles. The goal of this project is to understand the molecular regulation of silicification processes in natural diatom populations to better understand the processes controlling diatom productivity in the sea. Through culture studies and two research cruises, this research will couple classical measurements of silicon uptake and silica production with molecular and biochemical analyses of Silicification-Related Gene (SiRG) and protein expression. The proposed cruise track off the West Coast of the US will target gradients in Si and iron (Fe) concentrations with the following goals: 1) Characterize the expression pattern of SiRGs, 2) Correlate SiRG expression patterns to Si concentrations, silicon uptake kinetics, and silica production rates, 3) Develop a method to normalize uptake kinetics and silica production to SiRG expression levels as a more accurate measure of diatom activity and growth, 4) Characterize the diel periodicity of silica production and SiRG expression.

It is estimated that diatoms process 240 Teramoles of biogenic silica each year and that each molecule of silicon is cycled through a diatom 39 times before being exported to the deep ocean. Decades of oceanographic and field research have provided detailed insight into the dynamics of silicon uptake and silica production in natural populations, but a molecular understanding of the factors that influence silicification

processes is required for further understanding the regulation of silicon and carbon fluxes in the ocean. Characterizing the genetic potential for silicification will provide new information on the factors that regulate the distribution of diatoms and influence in situ rates of silicon uptake and silica production. This research is expected to provide significant information about the molecular regulation of silicification in natural populations and the physiological basis of Si limitation in the sea.

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## Funding

Funding Source	Award
<a href="#">NSF Division of Ocean Sciences (NSF OCE)</a>	<a href="#">OCE-1333929</a>
<a href="#">NSF Division of Ocean Sciences (NSF OCE)</a>	<a href="#">OCE-1334387</a>

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