

Results from experiments examining the cell diameter of *Crocospaera watsonii* (WH0003) in response to light irradiance; conducted in the Hutchins Laboratory, USC

Website: <https://www.bco-dmo.org/dataset/4066>

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Project

» [CO2 control of oceanic nitrogen fixation and carbon flow through diazotrophs](#) (Diaz N2-Fix in High CO2)

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Table of Contents

- [Dataset Description](#)
 - [Methods & Sampling](#)
 - [Data Processing Description](#)
- [Data Files](#)
- [Parameters](#)
- [Instruments](#)
- [Deployments](#)
- [Project Information](#)
- [Funding](#)

Dataset Description

Results of a laboratory experiment examining growth of the WH0003 isolate of *Crocospaera watsonii* in response to different light intensities. WH0003 was isolated near Sta. ALOHA (A Long Term Oligotrophic Habitat Assessment) in the North Pacific Ocean near Hawaii (22 deg 45' N, 158 deg 00' W).

Detailed methods and results are described in the following publication (see Figure 4, panel a): Garcia, N.S., Fu, F.X., and Hutchins, D.A. (2013). Colimitation of the unicellular photosynthetic diazotroph *Crocospaera watsonii* by phosphorus, light, and carbon dioxide. *Limnology and Oceanography* 58(4): 1501-1512. DOI: [10.4319/l.2013.58.4.1501](https://doi.org/10.4319/l.2013.58.4.1501)

Methods & Sampling

Culturing and experimental conditions

Experimental cultures were grown with a semi-continuous culturing method at 28 degrees C in autoclave-sterilized artificial seawater medium with nutrients added in concentrations equivalent to the recipe for the Aquil medium (except for NO₃⁻), as in Garcia et al. (2011) and originally described by Morel et al. (1979).

Light experiment and cellular growth rates

Triplicate cultures were grown in 800 mL polystyrene flasks under 5 irradiances (18, 40, 100, 180, 300 $\mu\text{mol quanta per m}^2 \text{ per second}$) and diluted every 2-3 days to 10-20 x 10³ cells per mL. Cells were counted microscopically in each replicate culture with a hemocytometer at the end of each dilution period, and steady state growth rates were calculated from an increase in culture cell number per unit volume between 2-3 dilution periods (4-6 days) after cultures were acclimated to treatment conditions for 7-10 generations. To calculate growth rates, the investigators used the equation $N_T = N_0 e^{\mu T}$, where N_0 and N_T are the initial and final culture cell densities, respectively and T is the amount of time in days between culture cell number estimates. With this method, the dilution rate is determined by the growth rate of the algae as determined by the

experimental treatments, rather than by controlling the growth rate through imposing a dilution rate, as one does for continuous cultures.

Cell diameters of ~12 cells from treatment replicates were measured with an ocular micrometer. In the light experiment, cells in one replicate from each light were measured treatment twice, once in the middle of the light period and once at the end of the light period on the same day.

Light was supplied on a 12:12 light:dark cycle with cool white fluorescent bulbs. The investigators terminally sampled each replicate culture 24 hours after the last dilution for N₂-fixation rates and CO₂-fixation rates.

References:

Garcia, N. S., F.-X. Fu, , C. L. Breene, P. W. Bernhardt, M. R. Mulholland, J. A. Sohm, and D. A. Hutchins. 2011. Interactive effects of irradiance and CO₂ on CO₂- and N₂ fixation in the diazotroph *Trichodesmium erythraeum* (Cyanobacteria). *J. Phycol.* 47: 1292-1303. DOI: [10.1111/j.1529-8817.2011.01078.x](https://doi.org/10.1111/j.1529-8817.2011.01078.x)

Morel, F. M. M., J. G. Rueter, D. M. Anderson, and Guillard, R. R. L. 1979. Aquil: Chemically defined phytoplankton culture medium for trace metal studies. *J. Phycol.* 15:135-141. DOI: [10.1111/j.1529-8817.1979.tb02976.x](https://doi.org/10.1111/j.1529-8817.1979.tb02976.x)

Data Processing Description

BCO-DMO re-arranged data formatted as separate tables into one dataset. Parameter names were changed to conform with BCO-DMO conventions.

[[table of contents](#) | [back to top](#)]

Data Files

File
C_watsonii_WH0003_cell_diam.csv (Comma Separated Values (.csv), 494 bytes) MD5:f655489fc74ef27e881c61de80cec831
Primary data file for dataset ID 4066

[[table of contents](#) | [back to top](#)]

Parameters

Parameter	Description	Units
time_period	In the light experiment, cells in one replicate from each light treatment were measured twice: once in the middle of the light (mid_light_period) period and once at the end of the light period (end_light_period) on the same day.	text
light	Light intensity. (For more about light measurement see: Australian National Algae Culture Collection and Plant Physiology Online.)	micromoles quanta per square meter per second (umol quanta m ⁻² s ⁻¹)
cell_diameter	Cell diameter measured in micrometers (um).	micrometers (um)
cell_diameter_se	Standard error of cell_diameter.	micrometers (um)

[[table of contents](#) | [back to top](#)]

Instruments

Dataset-specific Instrument Name	Hemocytometer
Generic Instrument Name	Hemocytometer
Dataset-specific Description	Cells were counted microscopically in each replicate culture with a hemocytometer at the end of each dilution period.
Generic Instrument Description	A hemocytometer is a small glass chamber, resembling a thick microscope slide, used for determining the number of cells per unit volume of a suspension. Originally used for performing blood cell counts, a hemocytometer can be used to count a variety of cell types in the laboratory. Also spelled as "haemocytometer". Description from: http://hlsweb.dmu.ac.uk/ahs/elearning/RITA/Haem1/Haem1.html .

Dataset-specific Instrument Name	Microscope-Optical
Generic Instrument Name	Microscope - Optical
Dataset-specific Description	Cell diameters were measured with an ocular micrometer.
Generic Instrument Description	Instruments that generate enlarged images of samples using the phenomena of reflection and absorption of visible light. Includes conventional and inverted instruments. Also called a "light microscope".

[[table of contents](#) | [back to top](#)]

Deployments

lab_Hutchins_07-12_diazotrophs

Website	https://www.bco-dmo.org/deployment/59043
Platform	USC
Description	Laboratory experiments conducted as part of project titled, "CO2 control of oceanic nitrogen fixation and carbon flow through diazotrophs".

[[table of contents](#) | [back to top](#)]

Project Information

CO2 control of oceanic nitrogen fixation and carbon flow through diazotrophs (Diaz N2-Fix in High CO2)

Coverage: Laboratory

From NSF award abstract:

The importance of marine N2 fixation to present ocean productivity and global nutrient and carbon

biogeochemistry is now universally recognized. Marine N₂ fixation rates and oceanic N inventories are also thought to have varied over geological time due to climate variability and change. However, almost nothing is known about the responses of dominant N₂ fixers in the ocean such as *Trichodesmium* and unicellular N₂ fixing cyanobacteria to past, present and future global atmospheric CO₂ regimes. Our preliminary data demonstrate that N₂ and CO₂ fixation rates, growth rates, and elemental ratios of Atlantic and Pacific *Trichodesmium* isolates are controlled by the ambient CO₂ concentration at which they are grown. At projected year 2100 pCO₂ (750 ppm), N₂ fixation rates of both strains increased 35-100%, with simultaneous increases in C fixation rates and cellular N:P and C:P ratios. Surprisingly, these increases in N₂ and C fixation due to elevated CO₂ were of similar relative magnitude regardless of the growth temperature or P availability. Thus, the influence of CO₂ appears to be independent of other common growth-limiting factors. Equally important, *Trichodesmium* growth and N₂ fixation were completely halted at low pCO₂ levels (150 ppm), suggesting that diazotrophy by this genus may have been marginal at best at last glacial maximum pCO₂ levels of ~190 ppm. Genetic evidence indicates that *Trichodesmium* diazotrophy is subject to CO₂ control because this cyanobacterium lacks high-affinity dissolved inorganic carbon transport capabilities. These findings may force a re-evaluation of the hypothesized role of past marine N₂ fixation in glacial/interglacial climate changes, as well as consideration of the potential for increased ocean diazotrophy and altered nutrient and carbon cycling in the future high-CO₂ ocean.

We propose an interdisciplinary project to examine the relationship between ocean N₂ fixing cyanobacteria and changing pCO₂. A combined field and laboratory approach will incorporate in situ measurements with experimental manipulations using natural and cultured populations of *Trichodesmium* and unicellular N₂ fixers over range of pCO₂ spanning glacial era to future concentrations (150-1500 ppm). We will also examine how effects of pCO₂ on N₂ and C fixation and elemental stoichiometry are moderated by the availability of other potentially growth-limiting variables such as Fe, P, temperature, and light. We plan to obtain a detailed picture of the full range of responses of important oceanic diazotrophs to changing pCO₂, including growth rates, N₂ and CO₂ fixation, cellular elemental ratios, fixed N release, photosynthetic physiology, and expression of key genes involved in carbon and nitrogen acquisition at both the transcript and protein level.

This research has the potential to evolutionize our understanding of controls on N₂ fixation in the ocean. Many of our current ideas about the interactions between oceanic N₂ fixation, atmospheric CO₂, nutrient biogeochemistry, ocean productivity, and global climate change may need revision to take into account previously unrecognized feedback mechanisms between atmospheric composition and diazotrophs. Our findings could thus have major implications for human society, and its increasing dependence on ocean resources in an uncertain future. This project will take the first vital steps towards understanding how a biogeochemically-critical process, the fixation of N₂ in the ocean, may respond to our rapidly changing world during the century to come.

[[table of contents](#) | [back to top](#)]

Funding

Funding Source	Award
NSF Division of Ocean Sciences (NSF OCE)	OCE-0722337

[[table of contents](#) | [back to top](#)]