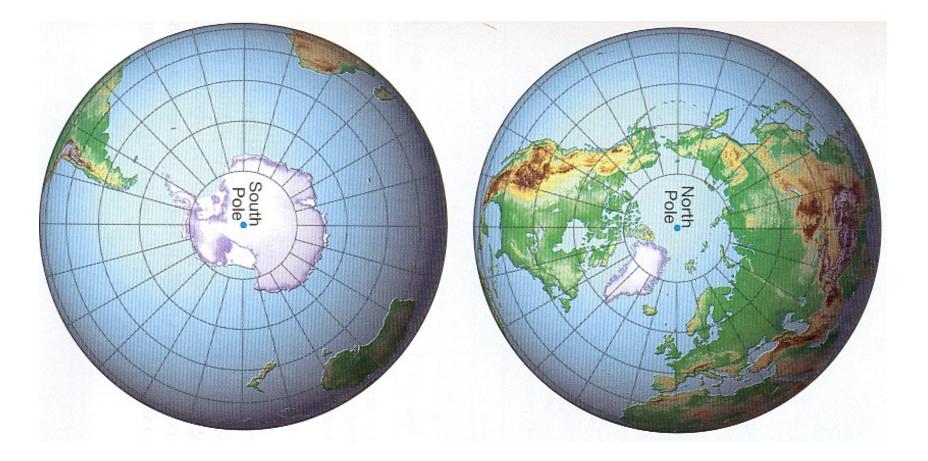
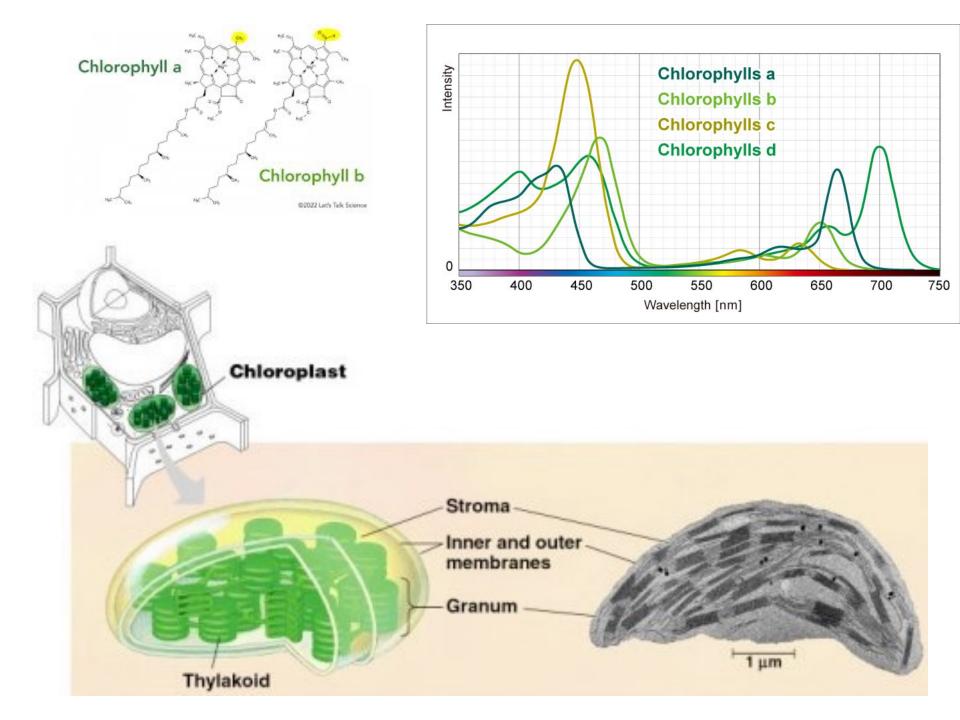
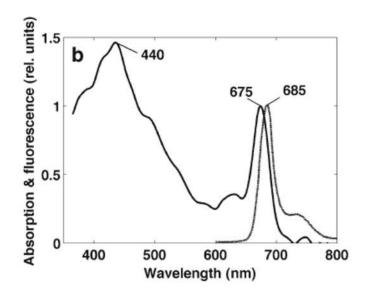
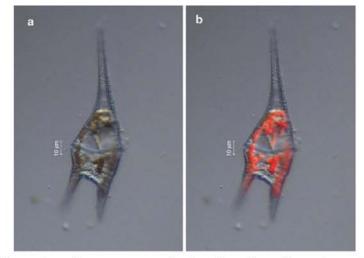
Description, techniques, protocols, and methodologies of key variables: chla & CDOM



Aurea Maria Ciotti – <u>ciotti@usp.br</u> Centro de Biologia Marinha da USP (<u>CEBIMar</u>)

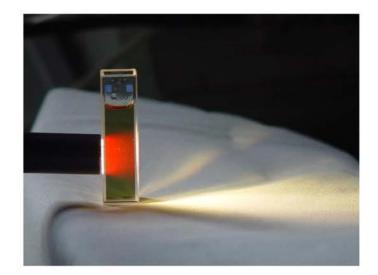






Chloroplast fluorescence in the dinoflagellate *Ceratium sp.* (*Photo: L.* Novoveska – Suggett et al, 2010)

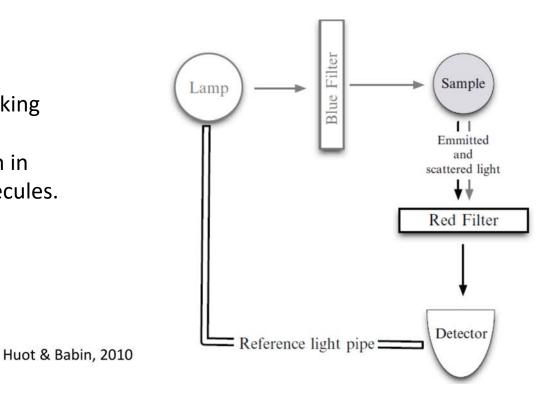
Fluorescence – one of the 3 main "fates" of light absorption by organisms that have photosystems



Fluorescence – easy, robust, sensitive – allowed to measure biology in the same space scales than Temperature in the ocean - "little green spheres"



- b) Photochemistry making more cells
- c) Fluorescence return in state for excited molecules.





Spectrophotometer





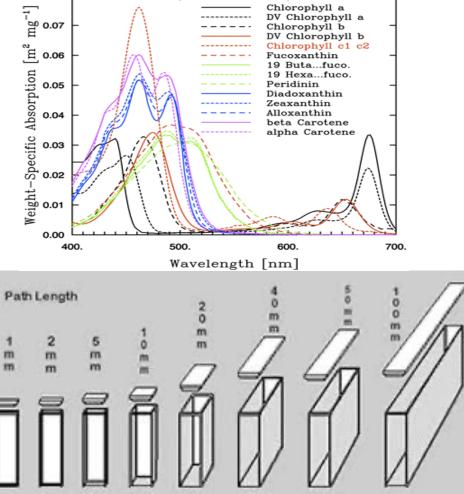
Fluorimeter



HPLC – High Performance Liquid Chromatography

Spectrophotometer

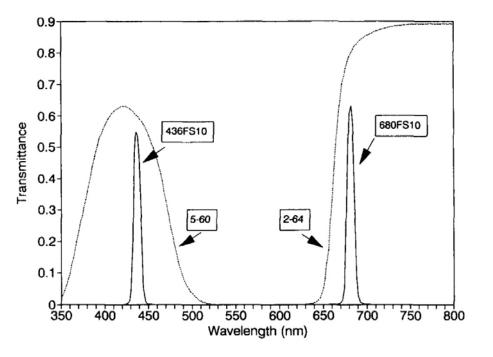




0.08

- a) Solvents and cuvette material
- b) Lamps and cuvette material

- a) Cleaning of material do not use HCl or after use soap and ultrapure water
- b) Try filtering samples as soon as possible grazing and degradation
- c) Avoid direct light and changes in temperature "room temperature"
- d) Emissions and excitations for fluorimeter



Welschmeyer DOI:10.4319/lo.1994.39.8.1985

Easy to calibrate – easy to verify drifts - solid standards and air measurements time series

10-AU fluorometer



Fluorometer Trilogy – Turner Designs

Aquafluor – hand held







Cyclops – Turner Designs



Fluorescence – in vitro; in vivo (excited by a known wavelength artificially of by sunlight, and variable/active

FRRf – fast repetition rate fluorometer



Azul ou multicolor



Phyto-PAM

Azul, vermelho ou multicolor

FIRe in situ

FIRe – Fluorescence Induction and Relaxation System

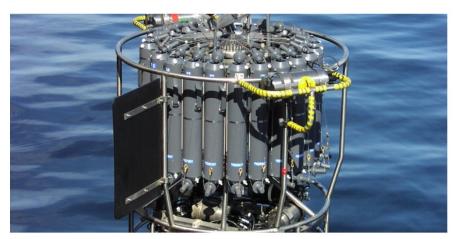


https://scor-int.org/group/156/

•Tortell, P.D. and Suggett, D.J. (eds) (2021) A User Guide for the **Application of Single Turnover** Active Chlorophyll Fluorescence for Phytoplankton Productivity Measurements. Version 1. Scientific Committee on Oceanic Research Working Group 156, 20pp. DOI: http://dx.doi.org/10.25607/OBP-1084

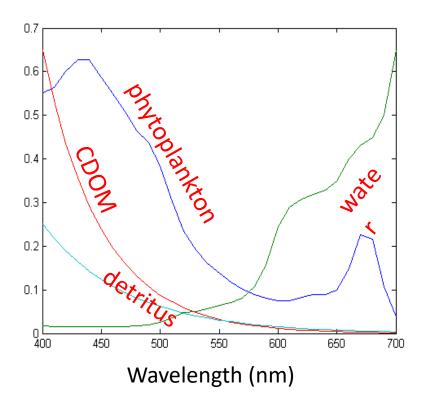
SCOR Working Group 156

Active Chlorophyll fluorescence for autonomous measurements of global marine primary productivity



Spectral Shapes for Optical active Components

 $a_t(\lambda) = a_w(\lambda) + a_{ph}(\lambda) + a_g(\lambda) + a_d(\lambda)$ total water phyto- CDOM detrital plankton particulate



CDOM and Chl compete for blue light

CDOM and the detrital material are very hard to discriminate, but globally a small part of at(l) budget – not the case for Rio de La Plata

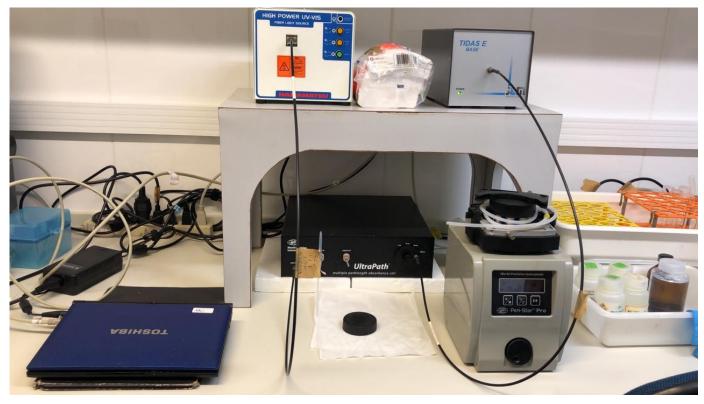
- Colored Dissolved Organic Matter (CDOM)
- gelbstoff, gilvin, yellow matter, chromophoric DOM
- a small fraction of the total DOM, variable composition
- CDOM is defined operationally by filtering Using a 0.2 μ m filter Absorption relative to "pure water" standard
- Typically, spectrum decreases exponentially $a_g(\lambda) = a_g(\lambda_o) \exp(-S(\lambda - \lambda_o))$

S varies from 0.015 to 0.024 $nm^{\text{-}1}$

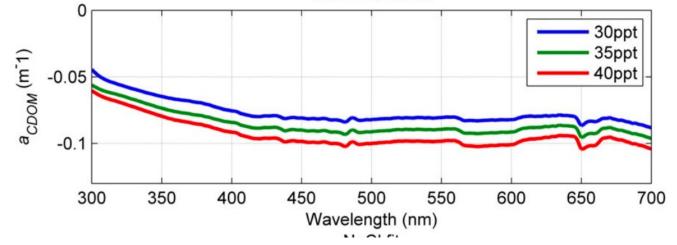
- Colored Dissolved Organic Matter (CDOM)
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NaCl measured



Pathlength in important

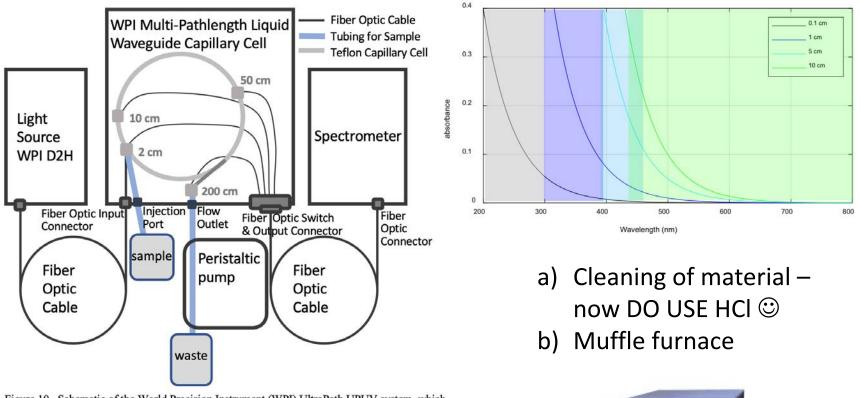
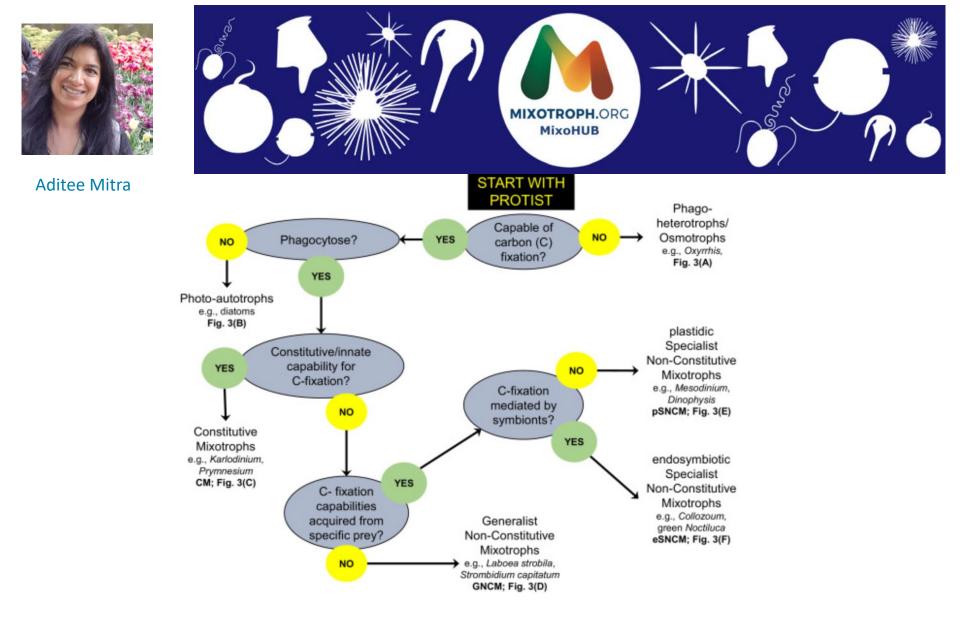


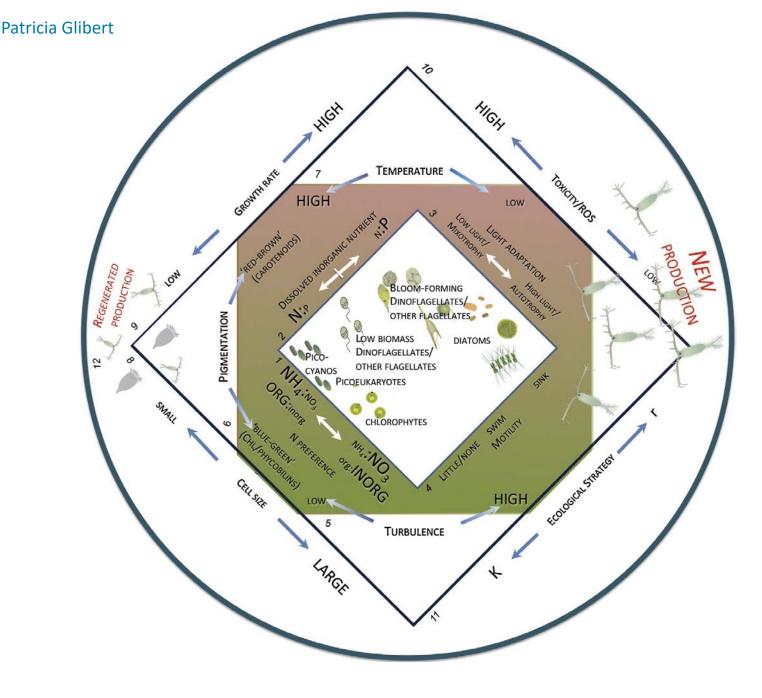
Figure 10. Schematic of the World Precision Instrument (WPI) UltraPath UPUV system, which includes the UltraPath absorbance sample cell (center) with four nominal optical pathlengths (2, 10, 50 and 200 cm), deuterium/tungsten light source (left), photodiode array spectrometer (right), and peristaltic pump.

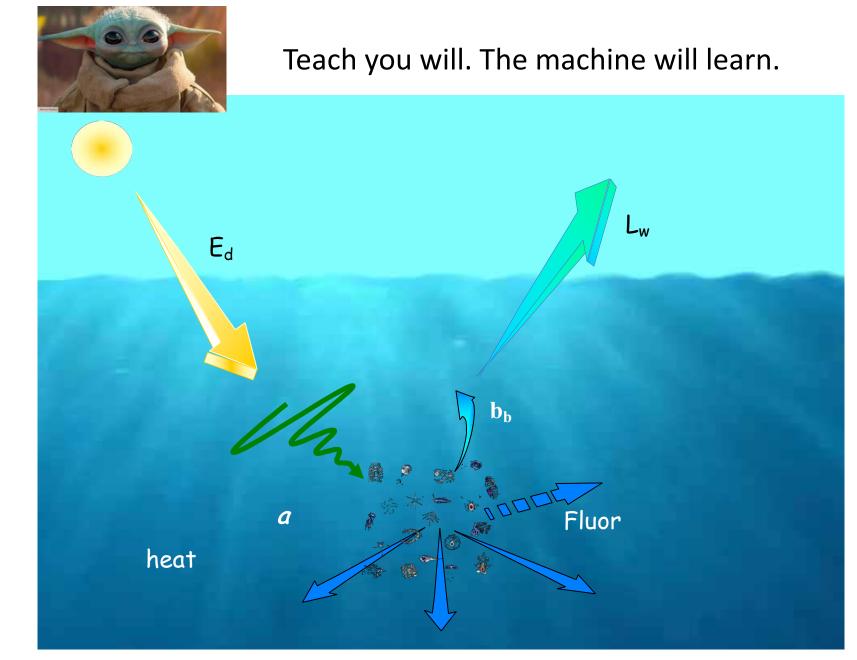




Mitra et al., 2016 10.1016/j.protis.2016.01.003







Yoda