

# OCB Theme II : Carbon uptake and storage

“Recent observational and modeling findings quantifying the magnitude and trends in ocean carbon fluxes and storage”

## **Autonomous measurements of the subpolar North Atlantic spring bloom: early results from the NAB08 experiment**

Mary Jane Perry  
University of Maine  
and NAB08 Colleagues

22 July 2008

## **Co-authors:**

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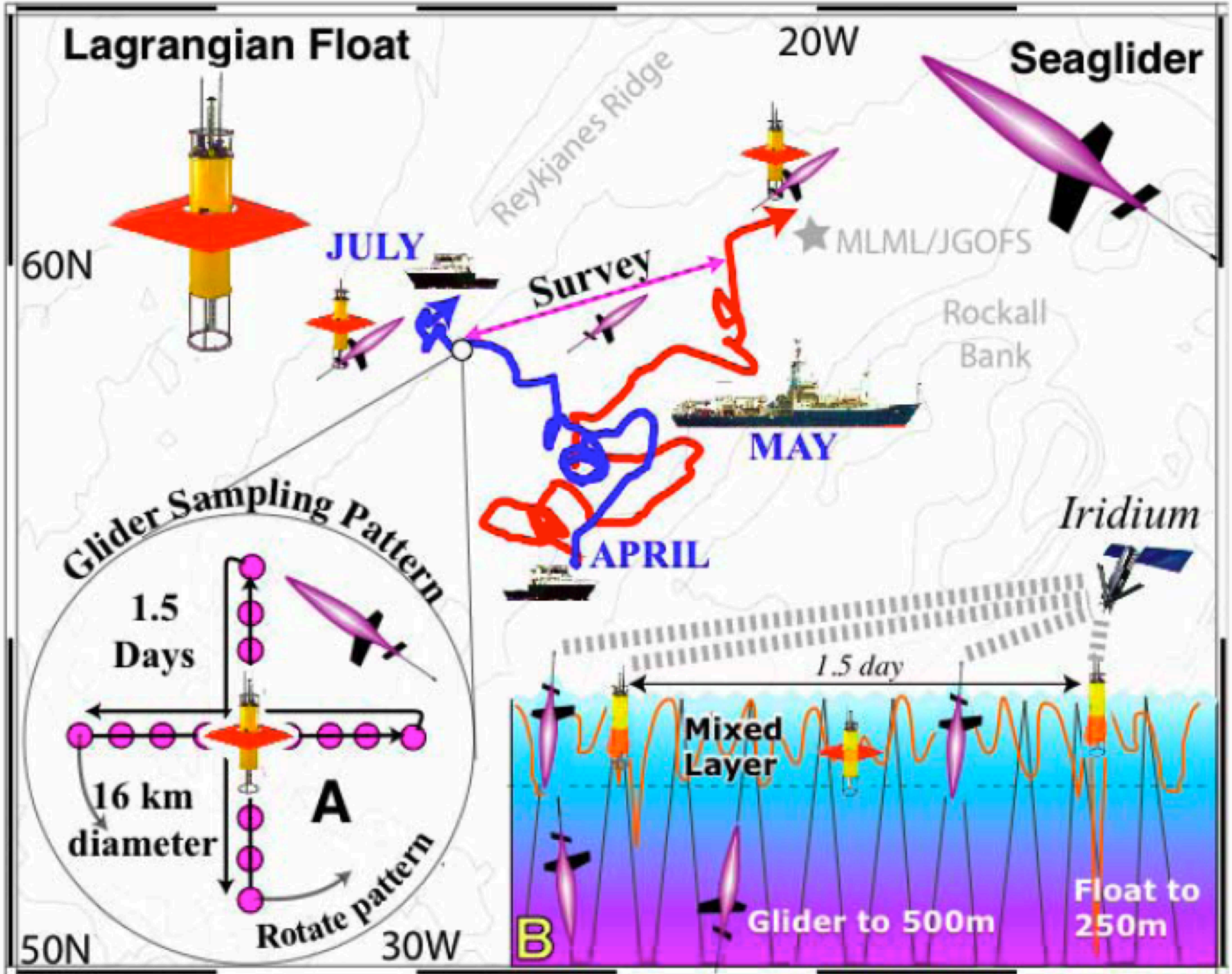
## **US (with support from NSF and NASA):**

UW, UMaine, Dalhousie, URI, Scripps, MBARI, OSU

## **International:**

Canada, Iceland, Denmark, UK (3 institutions)

**Thanks** to UW APL float and Seaglider groups for great jobs on floats 47, 48 and Seagliders 140, 141, 142, 143



## **So, why the subpolar North Atlantic ?**

### **Carbon uptake:**

One of largest CO<sub>2</sub> draw-downs on the planet occurs during NA spring bloom → photosynthetic uptake of carbon

### **Carbon Removal: 3 mechanisms**

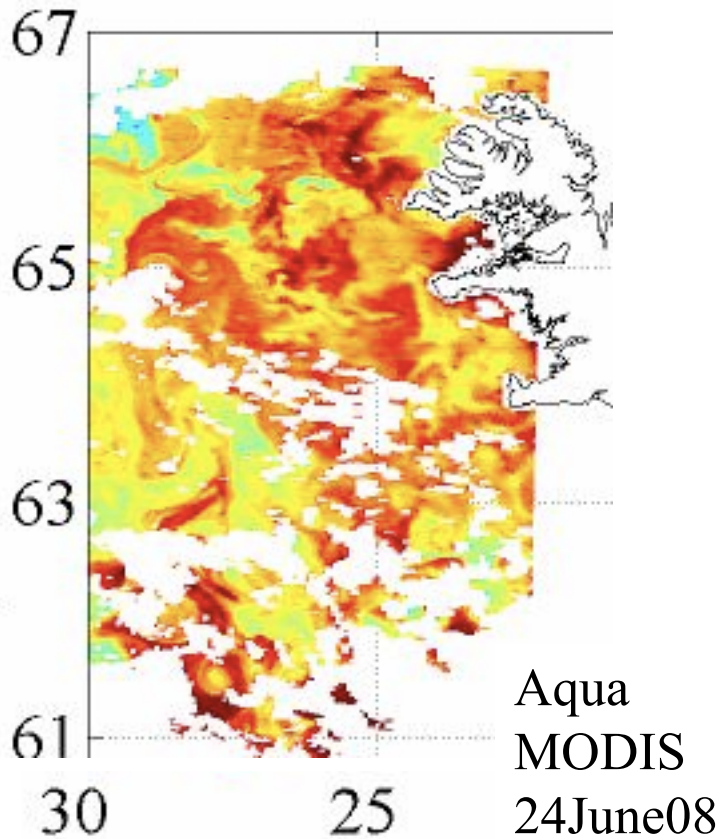
- \* mixed-layer pump (stratification/destratification)
- \* sinking of aggregates (diatoms, etc.)
- \* subduction of water mass

### **Carbon Storage:**

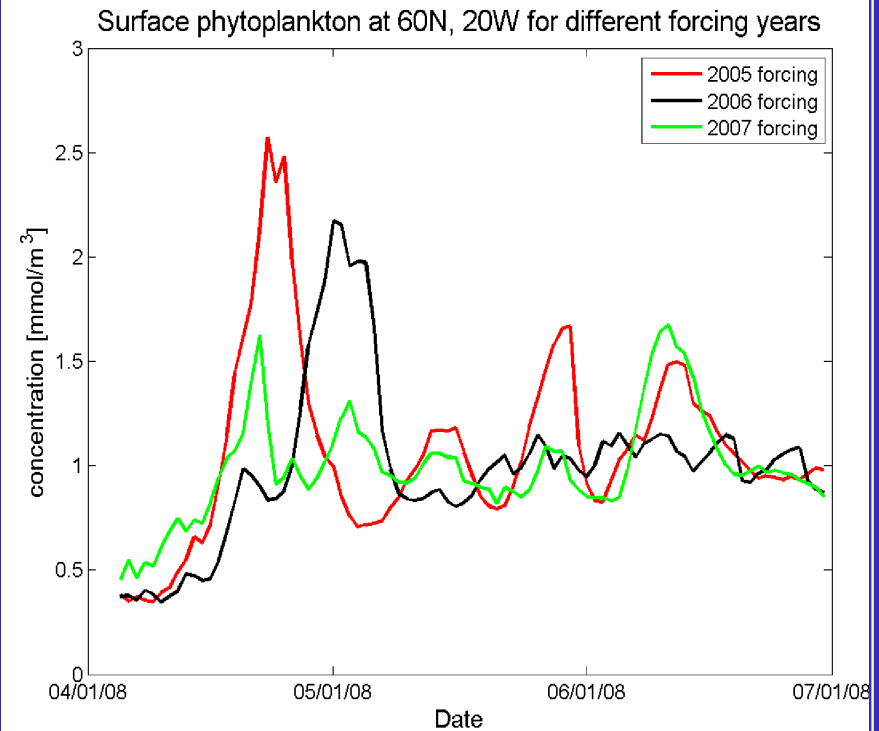
depends on – how much gets down, how fast, and what happens next . . .

# Challenges to studying NA bloom (and other regions)

High spatial variability on mesoscale and submesoscale (~ km)



Inter-annual variability in timing of bloom (1D model, similar to SeaWiFS data)



Fennel and Bagniewski

## Key OCB questions:

- 1) how much carbon is taken up ?
- 2) how much carbon is removed ?
- 3) how much carbon is stored ?
- 4) how will carbon uptake, removal and storage in the N. Atlantic, and other key regions, respond to **climate change or increased variability in forcings** ?
- 5) how can one **document change** – in light of **large temporal and spatial variability** ?

## Challenges to assessing carbon uptake and removal:

### **Ships –**

operate on fixed schedules (match or mismatch with timing of the bloom and major removal events)

### **Moorings –**

single locations, how to interpret submesoscale variability ?

### **Satellites –**

can't see through (persistent) clouds; lack depth resolution

### **Models –**

depend on quality of input data and understanding of processes

***The Project:*** “NSF-Collaborative Research: Autonomous Measurements of Carbon Fluxes in the North Atlantic Bloom”

New approach to studying the evolution and demise of the subpolar North Atlantic spring bloom near 60°N JGOFS site using floats and gliders, ship-based observations, satellites and models, and laboratory.

***The Motivation:***

- 1) JGOFS NABE synthesis and modeling activities clearly identified a need for **more complete temporal / spatial coverage of the bloom** and **improved resolution of mixed-layer dynamics and lateral processes**.
- 2) **Autonomous platforms now sufficiently mature** to carry out a three-month, open-ocean experiment.



## Core PIs, students and responsibilities

Eric D'Asaro, Eric Rehm

Katja Fennel, Witold Bagniewski  
(collaboration through student)

Craig Lee, Amanda Gray

Mary Jane Perry, N. Briggs, E. Kallin

Michael Sieracki, Nicole Poulton

Annette deCharon

Lagrangian bio-floats  
ecosystem models

Seagliders

ship optics & samples

phytoplankton species

education and outreach



**Process cruise collaborators**

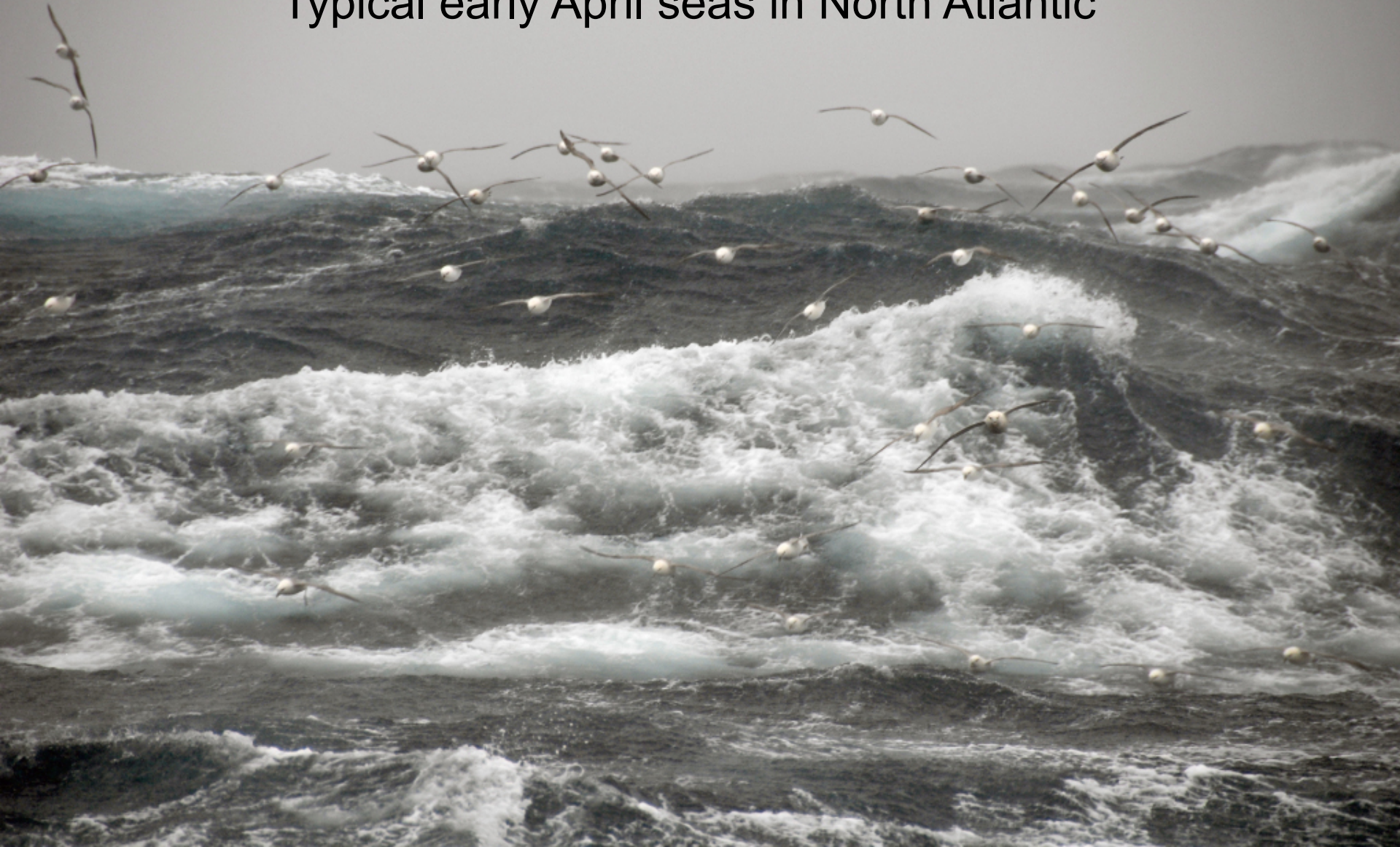
## Approach:

- \* autonomous 4-D sampling for 3 month, April–June
- \* 2 heavily-instrumented floats and 4 Seagliders
- \* proxy sensors for carbon-cycle components
- \* deploy all before spring bloom starts
- \* retrieve after spring bloom
- \* ship visits to help interpret data
  - \*\* ancillary measurements on 3-week process cruise with great input from collaborators (e.g., R. Lampitt's floating sediment traps)
- \* satellite data
- \* ecosystem model

# Deployment cruise: 1-6 April 2008



Typical early April seas in North Atlantic



# Two Lagrangian bio-heavy floats

water-following  
T, C (2 each)  
O<sub>2</sub> (2 types)  
Transmission (c)  
Chl fluorescence  
Backscatter (2 $\lambda$ )  
Ed ( $\lambda$ ) and Lu ( $\lambda$ )  
PAR  
ISUS NO<sub>3</sub><sup>-</sup>





## Four Seagliders

float-following

T, C

O<sub>2</sub> (2 types)

Chl fluorescence (2)

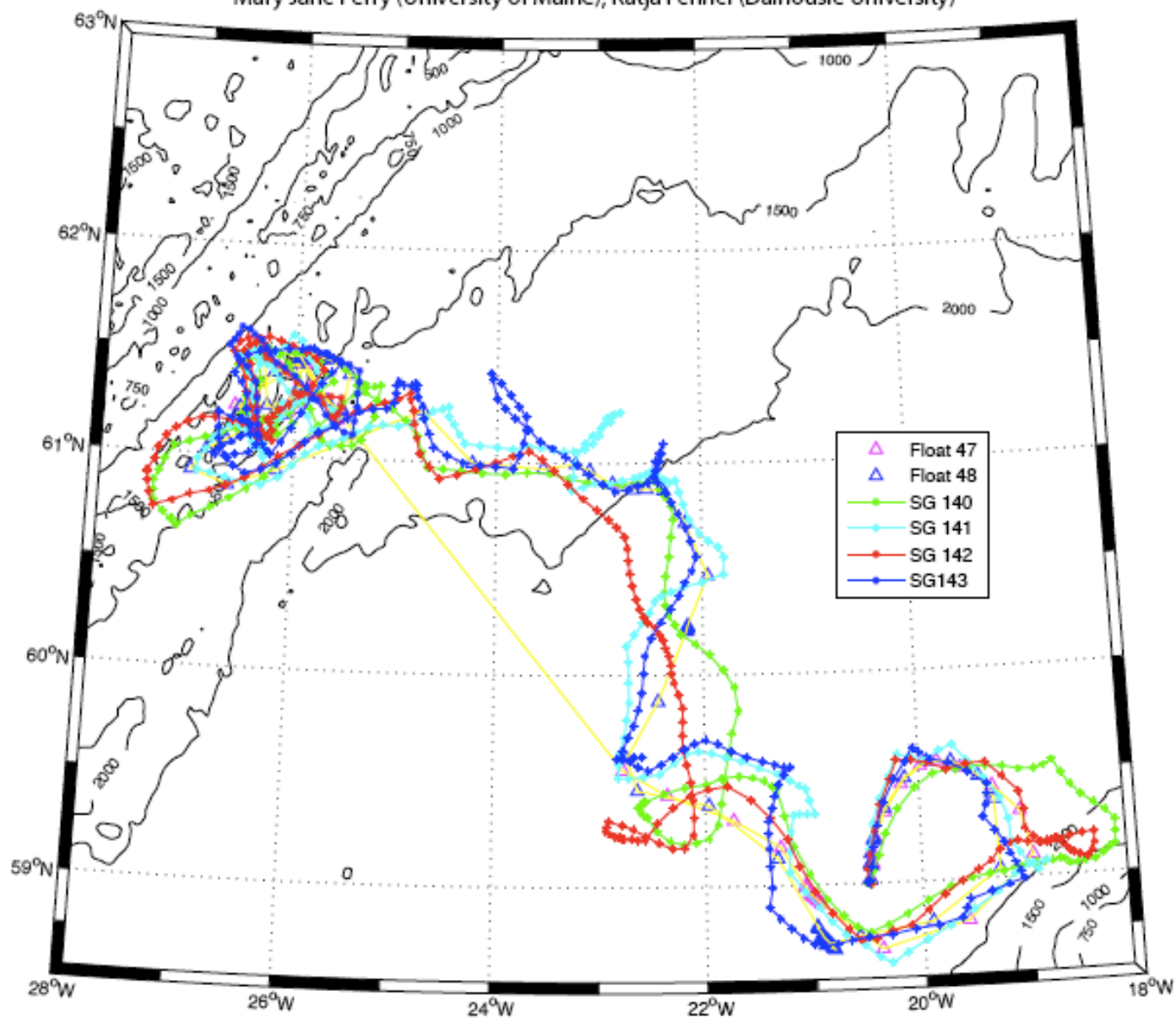
Backscatter (3 $\lambda$ )

CDOM fluorescence

# North Atlantic Bloom 2008: 4 April - 22 May 2008

Locations of Lagrangian Floats and Seagliders

PIs: Eric D'Asaro and Craig Lee (University of Washington - Applied Physics Lab),  
Mary Jane Perry (University of Maine), Katja Fennel (Dalhousie University)



**3 months of data, 4 Seagliders - deployed before bloom**

**May 2, ship arrived on station after beginning of bloom**

*Panels with backscattering, chlorophyll from  
fluorescence, dissolved oxygen,  
chlorophyll/backscattering*



# **Value of process cruise**

(some limited activities on deployment/rescue/pickup cruises)

**Sensor calibration** (vicarious)

**Validation** of 'proxy' relationships

**Enhanced interpretation**

of proxies and processes

# **“Calibration”**

## **Has the sensor drifted or fouled?**

Aggressive CTD calibration program

Cross-platform, vicarious sensor calibrations for:

temperature and transmission

conductivity

oxygen

chlorophyll fluorescence

backscattering

PAR and spectral (ir)radiance

# Float 48 calibration operation



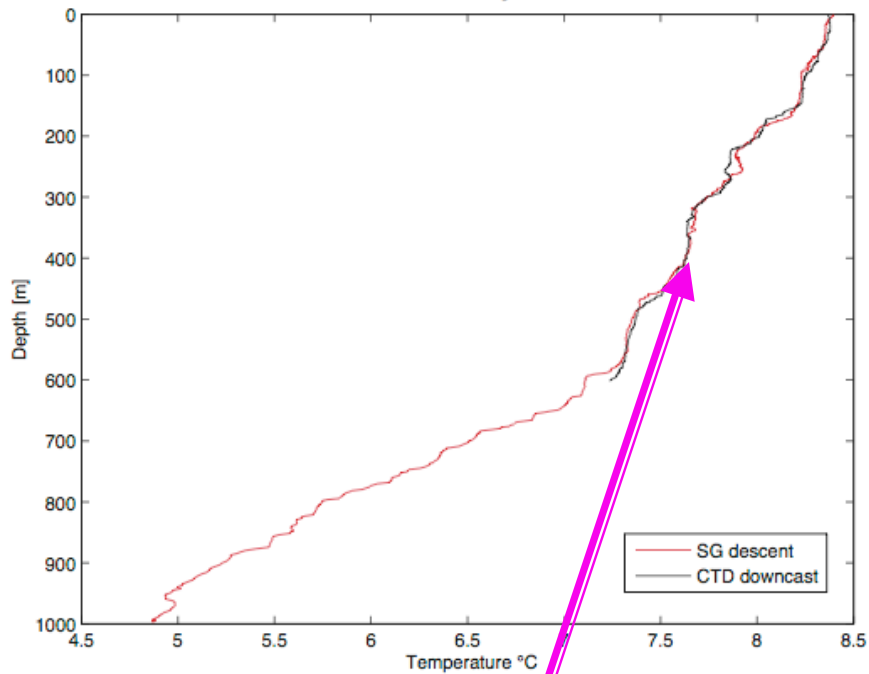
# Seaglider calibration operation



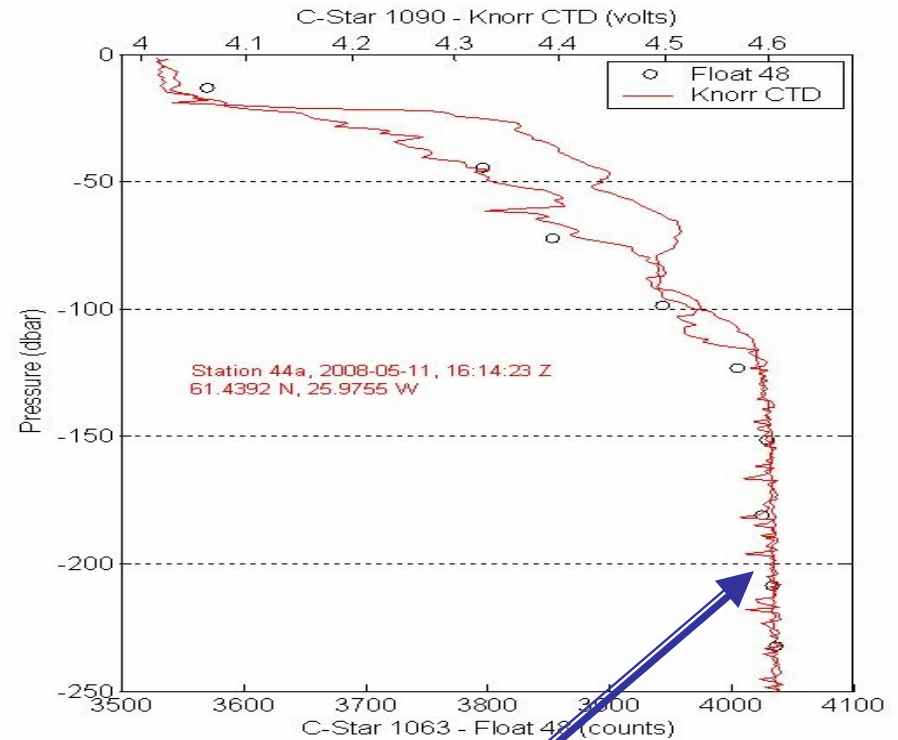
# “Calibration”

## Has the sensor drifted or fouled?

SG142 CTD Cal R/V Knorr: Temperature  
CTD Cast Station 12a: 06-May-2008 09:33 UTC  
SG Dive 151: 06-May-2008 09:40 UTC



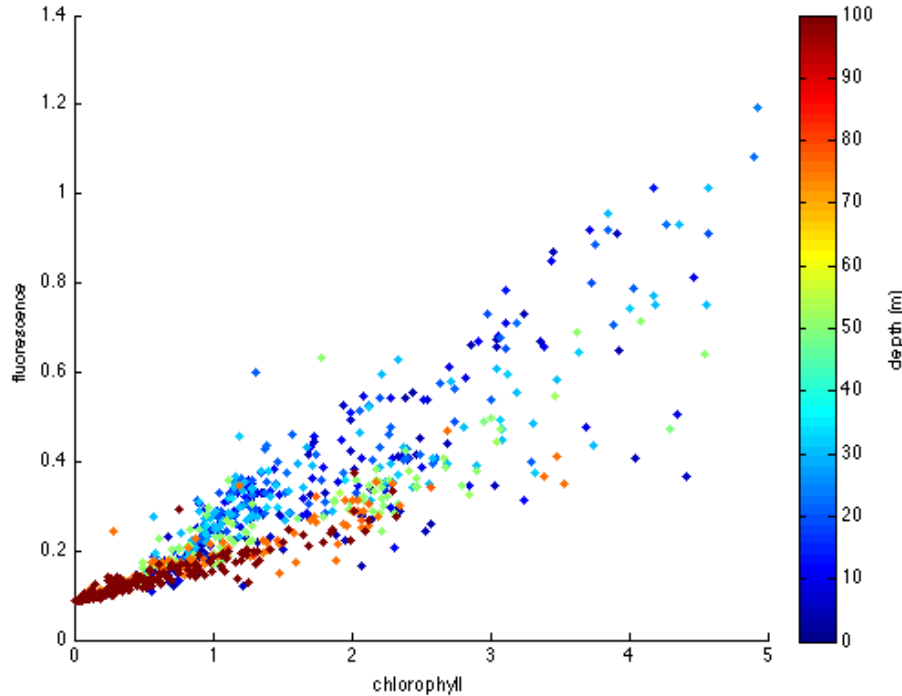
Temperature



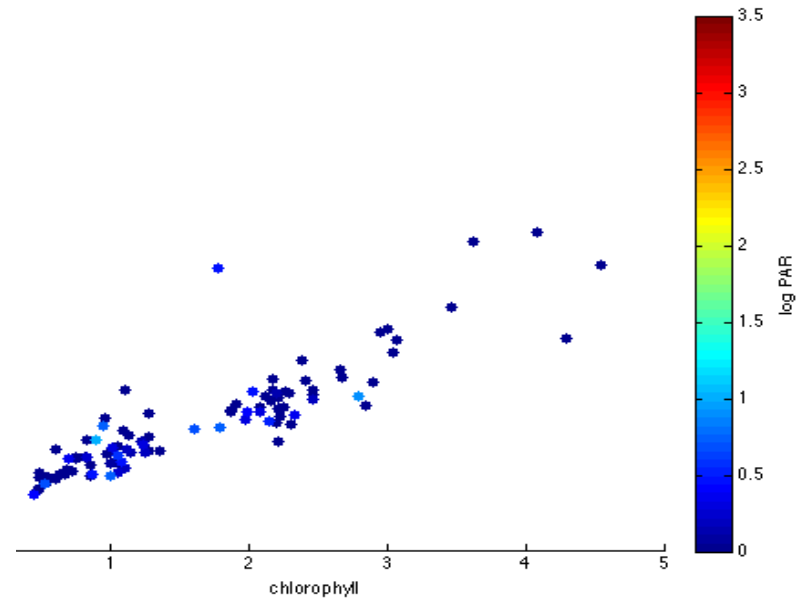
Transmission

# “Validation of Proxies”

chlorophyll fluorescence vs. extracted chlorophyll:  
fluorescence —> chlorophyll



all data (1,000 points)  
colored by depth



50-m data  
colored by PAR

# “Validation of Proxies”

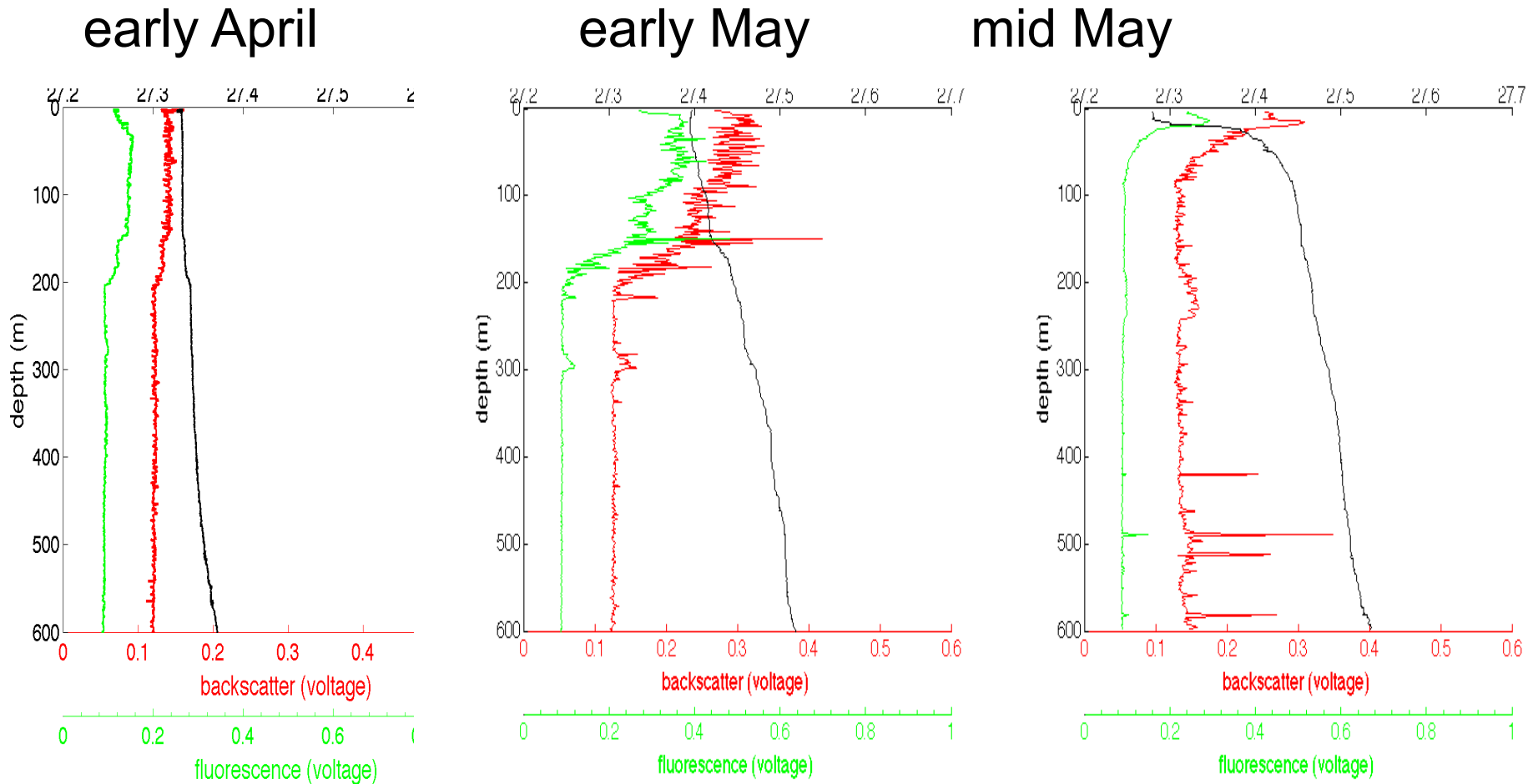
Other proxies remaining to be examined:

attenuation ( $c$ ) and $b_b$	vs.	<b>POC</b> (~1,000 samples & blanks)
ISUS <b>nitrate</b>	vs.	chemical measurement ~1,000 samples)
chlorophyll	vs.	<b>phyto carbon</b> from FCM, FlowCAM
$Lu(\lambda) / Ed(\lambda)$	vs.	<b>phytoplankton “groups”</b> from HPLC pigments & FCM, FlowCAM

# “Enhanced interpretation of optical & other signals”

## 1) Shift in phytoplankton community structure

Large diatoms chains produce high-frequency variability in surface chl fluorescence

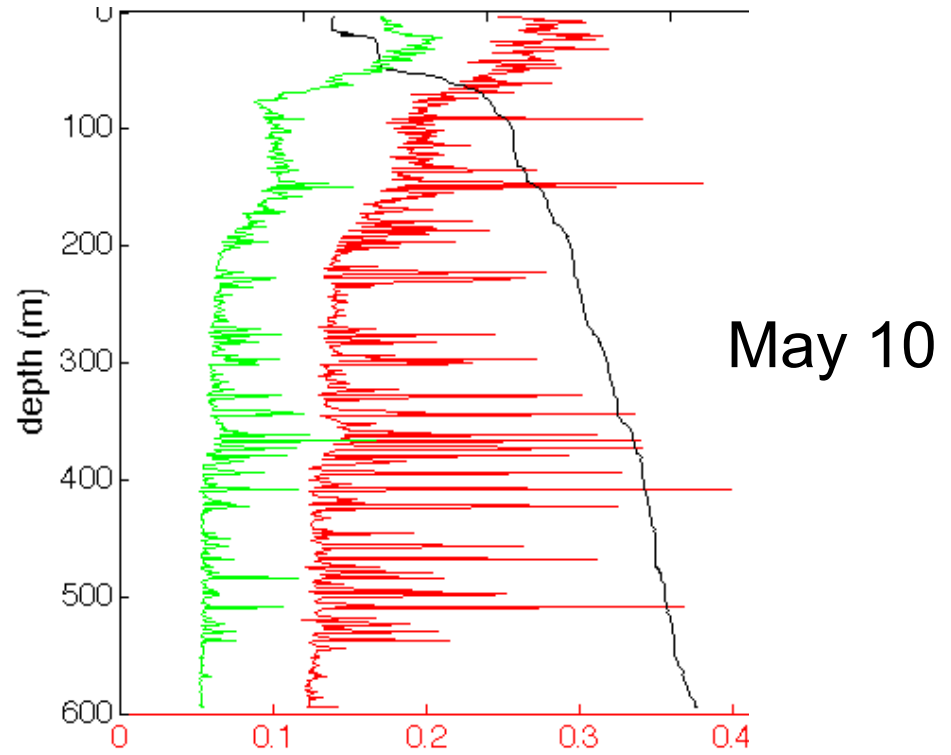
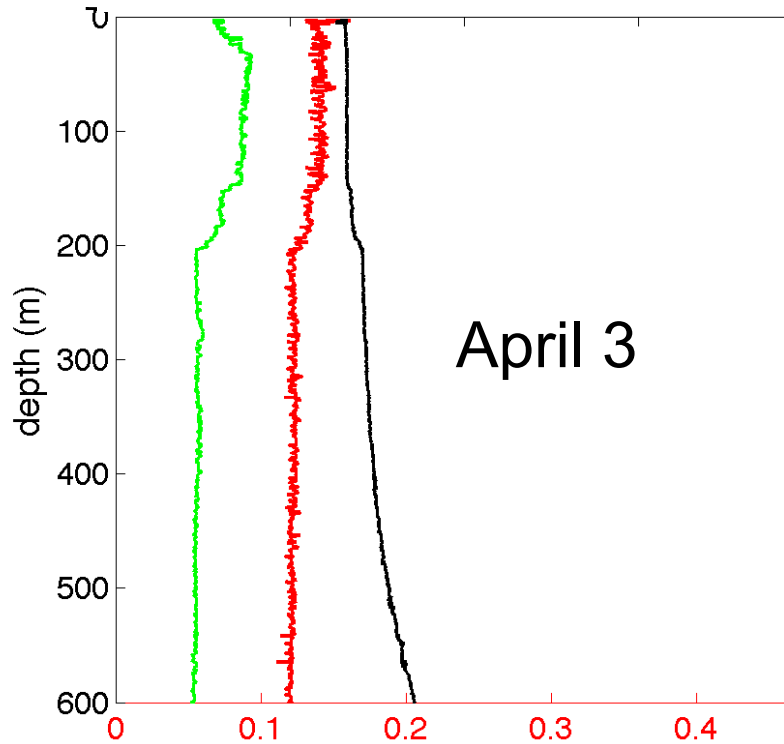




# “Enhanced interpretation of optical & other signals”

## 2) Carbon flux:

high-frequency variability in deep chlorophyll fluorescence and backscatter, AKA “spikes”



**red = bb; green = chl F; black = density**

Seaglider 142 backscatter, deepening horizon of particle spikes  $\sim 50 \text{ m d}^{-1}$

Large PELAGRA trap catches at  $\sim 600 - 700 \text{ m}$  coincide with backscatter events

What initiated the flux? Why so rapid? Probably silicate (not yet analyzed). ISUS data on float suggests nitrate was not depleted.

What comprised the sink? Large chain forming diatoms.