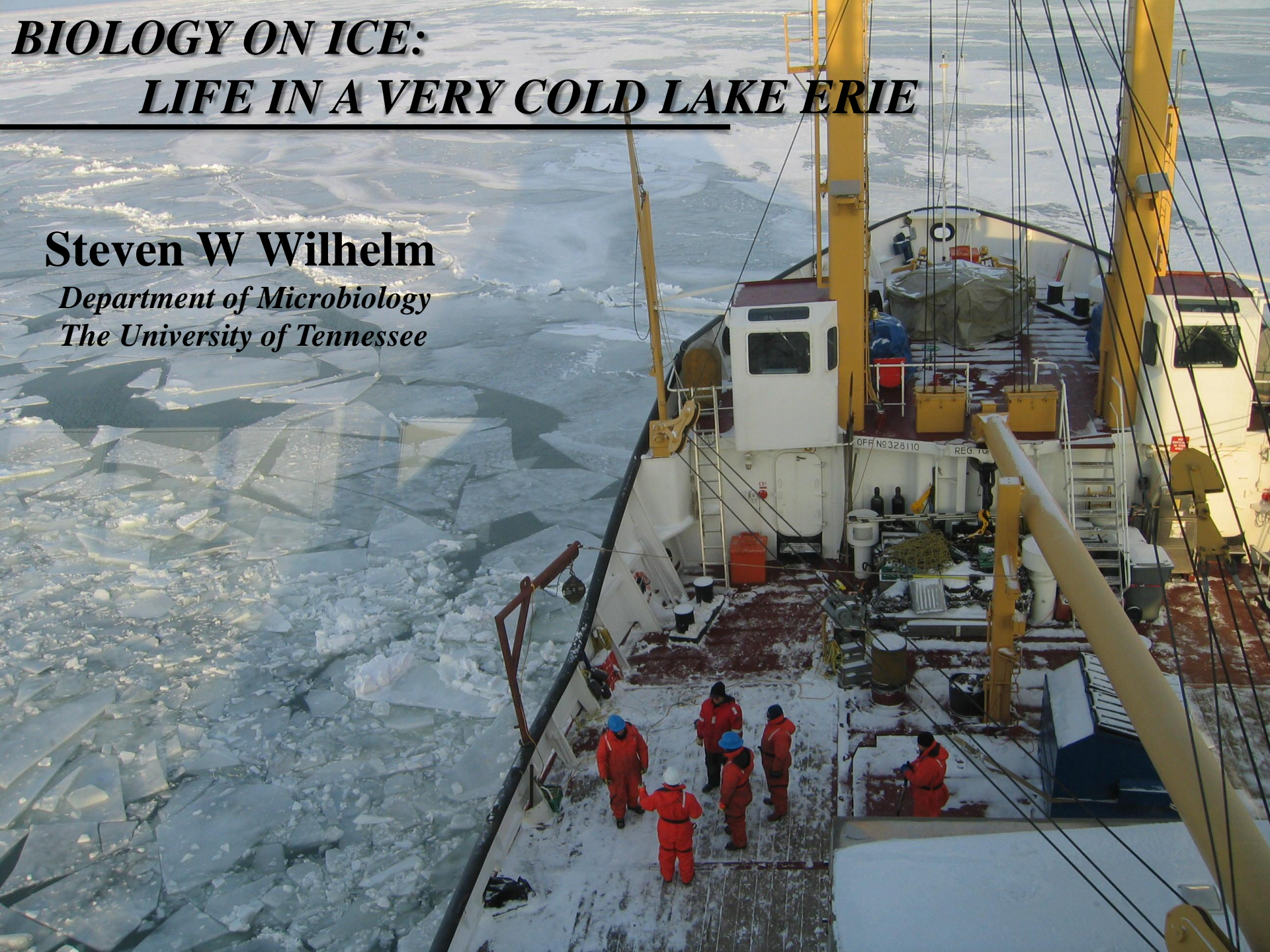


BIOLOGY ON ICE: LIFE IN A VERY COLD LAKE ERIE

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WINTER ASSESSMENT OF MICROBIAL BIOMASS AND METABOLISM

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With special thanks to

The Captains and crews of the

CCGS Griffon

Alice Dove

(Environment Canada)

Vi Richardson

(Environment Canada)

Steve Smith

(Environment Canada)

Todd Breedon

(Environment Canada)

Dave Gilroy

(Environment Canada)

SUPPORT FROM

Environment Canada

Canadian Coast Guard

Ohio Lake Erie Commission

Ohio Sea Grant

New York Sea Grant

Most photos from most Lake Erie cruises look like this one



To develop any forecasting power, we need to look beyond May - October

Life in (on, under) a very cold Lake Erie

A brief history of winter work on Lake Erie

(brief because that is all there is)

Winter Assessment of Biomass and Metabolism

WAMBaM, 2007 - 2010

- What have we been doing?***
- What do we know?***
- What does it all mean?***

Previous winter limnology on Lake Erie Last major study was seventy (70) years ago.

Chandler 1940, Chandler 1942a, Chandler 1942b, Chandler & Weeks 1945

THE OHIO JOURNAL OF SCIENCE

VOL. XL

NOVEMBER, 1940

No. 6

LIMNOLOGICAL STUDIES OF WESTERN LAKE ERIE

I. PLANKTON AND CERTAIN PHYSICAL-CHEMICAL DATA OF THE BASS ISLANDS REGION, FROM SEPTEMBER, 1938, TO NOVEMBER, 1939

DAVID C. CHANDLER

Franz Theodore Stone Laboratory
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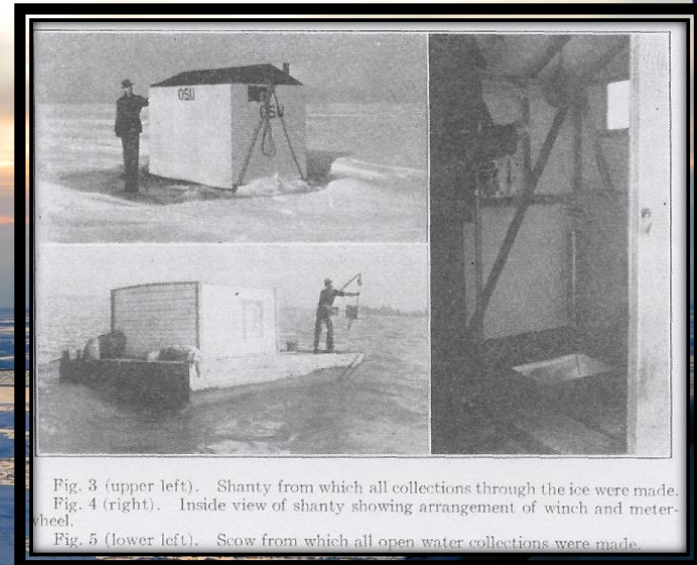


Fig. 3 (upper left). Shanty from which all collections through the ice were made.
Fig. 4 (right). Inside view of shanty showing arrangement of winch and meter-wheel.
Fig. 5 (lower left). Scow from which all open water collections were made.

Chandler observed “pulses” (blooms) of diatoms in mid-winter (mid February to late March) under the ice, low zooplankton populations, and a variable light environment.

A number of “snap shots” taken since then –
Rockwell, D.C., Salisbury, D.K., and Lesht, B.M. 1989.

few process measurements -
e.g., on Lake Michigan - Scavia and Laird 1987; Scavia et al. 1986

TIMELINE (What have we been doing). Assessment of biological limnology during the “winter”

**2007 – first WAMBaM survey (Wilhelm, McKay, Twiss, Bullerjahn – 2 day “ride along”)
- noted significant biomass in the lake**

**2008 – WAMBaM II. Measured significant rates of primary production. Repeated observations on biomass.
Collected samples for thorough community characterization and bacterial production estimates.**

**2009 - WAMBaM IIIa and IIIb. Measured significant rates of primary production. Repeated observations on
biomass. Collected samples for thorough community characterization and bacterial production estimates.
Surveys shortened due to logistics.**

2010 - WAMBaM IV. Survey shortened due to logistics. Most data still processing.

Research complimented by US Coast Guard (Neah Bay) and EPA (e.g., Lake Guardian) cruises



Is biomass in ice, under ice and/or in open water?

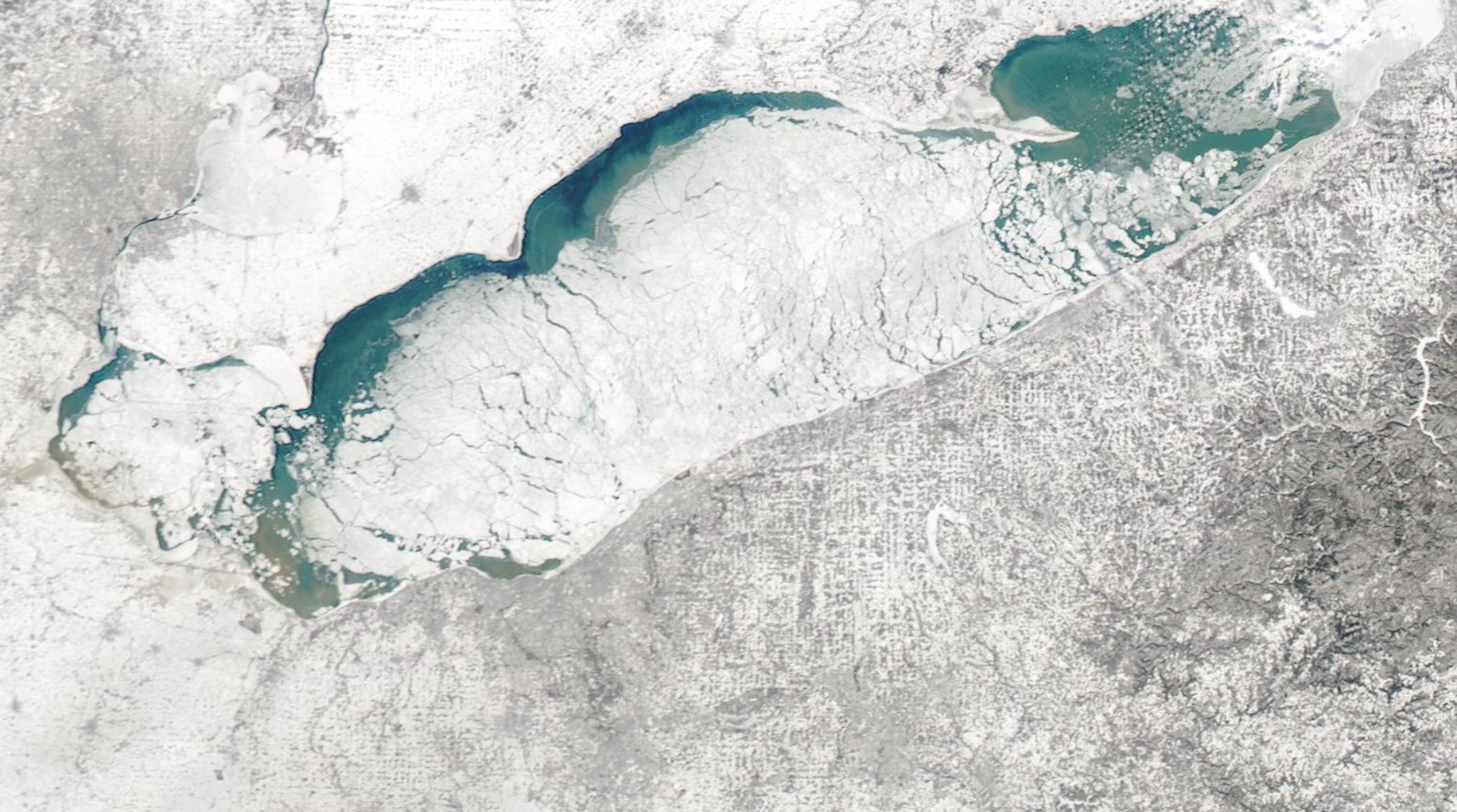
CACHE: Concentrated Algae Community and Heterotrophic Ecosystem

- **Extremely high chlorophyll**
- **Rapidly growing phytoplankton**
- **Abundant bacteria**



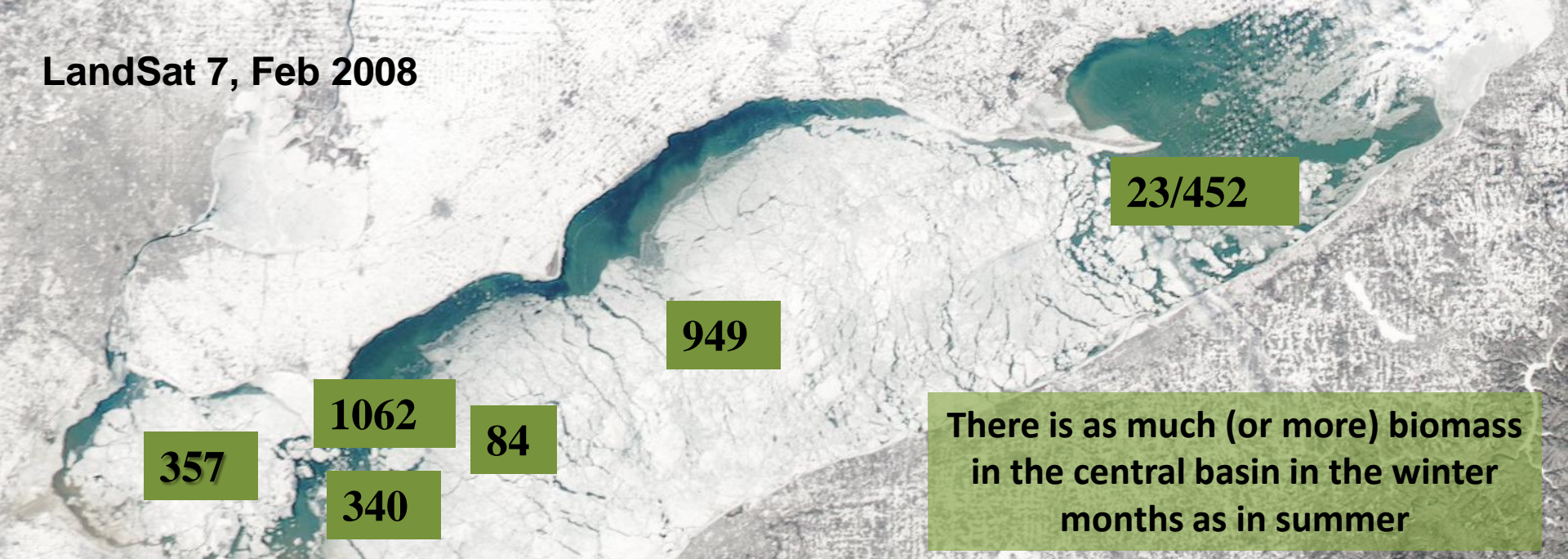
Is biomass in ice, under ice and/or in open water?





Ice is transient. It comes and goes. Commonly moves northwest to southeast (away from the shore) and southwest to northeast (with the wind)

LandSat 7, Feb 2008



Chlorophyll comparisons ($\mu\text{g L}^{-1}$)

Station	Feb 2007	Feb 2008	Feb 2009	Feb 2010	Jul 2005	Aug 2005	Aug 2006	Aug 2007
23/452	0.6	0.6	0.6	1.4	1.4	1.6	0.6	3.1
84	2.4	2.4	0.7	<u>1.4</u>	1.5	4.4	0.9	3.6
357	1.2	0.6	0.4	na	3.3	15.1	5.1	3.7
340/961/341	<u>8.4</u>	2.4	<u>9.66</u>	<u>1.4</u>	2.5	10.4	1.3	na
949/1053	na	1.5	2.21	2.1	na	na	na	na



ice-free winter samples

na – not available

All samples collected at 1m

CONSTRAINTS ON PRIMARY PRODUCTION

LOTS OF NUTRIENTS

Direct measures of water column chemistry as well as microcosm growth assays suggest the community is nutrient replete.

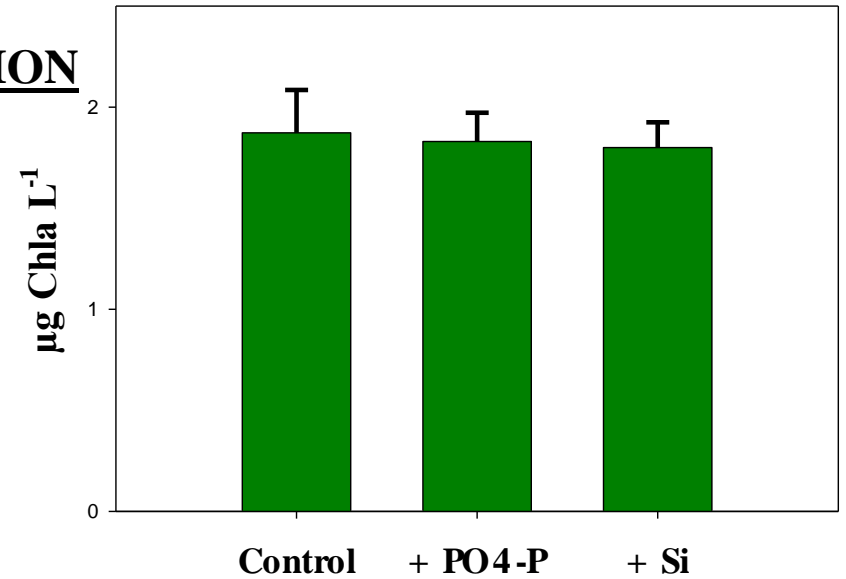
Alkaline phosphatase extremely low (not shown).

LIGHT

There is significant light penetration to at least 3 m in the water column and the community seems to be able to rapidly respond to levels of high light.

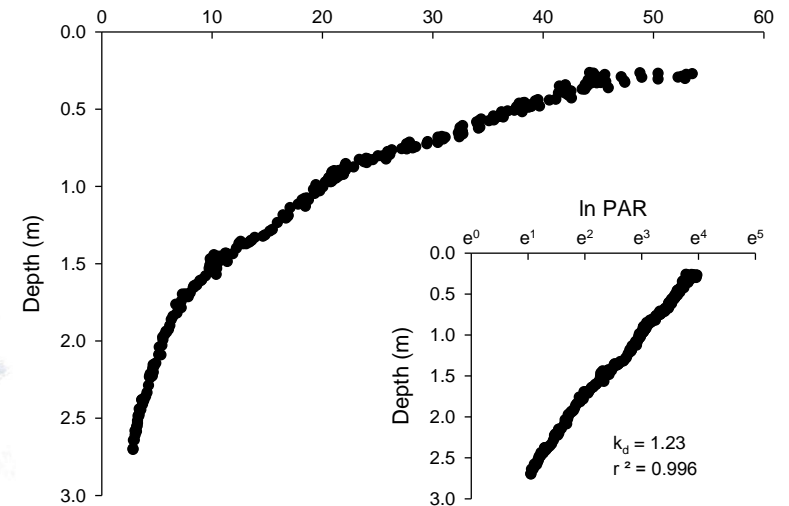
Light limitation seems likely though as the water column is isothermal.

Station 340 Growout



Wilhelm and Saxton 2008

Station 84 light penetration (PAR)



Harrison and Smith 2010

RESULTS - 2008: Total (>0.2 μm) primary production (data from MR Twiss) ranged from 0.7 to 1.7 g C·gChl- a^{-1} · h^{-1} , and are similar to those measured by McKay (^{14}C technique, photosynthetron) and Carrick (light:dark DO technique).

The concentration of picoplankton was too low at Sta 357 (3% total) to accurately measure ^{14}C uptake and accumulation.

Values are ca 20-60% of values measured using the same technique during summer 2003 and incubated at 150-300 $\mu\text{mol photons}\cdot\text{m}^{-2}\cdot\text{s}^{-1}$ (Ostrom et al. 2005). The higher winter biomass in some areas suggests total production is similar.

Station	Size fraction	Chlorophyll- <i>a</i>		Primary production
		$\mu\text{g Chl-}a/\text{L}$	%Total	gC/g Chl- <i>a</i> /hr
Mean \pm standard deviation ($n = 3$)				
357	Micro	2.41	66	0.74 ± 0.20
	Nano	1.10	30	0.85 ± 0.24
	Pico	0.12	3	-1.52 ± 1.73
	Total	3.63	---	0.70 ± 0.05
84	Micro	0.88	58	1.28 ± 0.22
	Nano	0.49	32	1.74 ± 0.28
	Pico	0.16	10	3.66 ± 1.56
	Total	1.53	---	1.68 ± 0.14
23	Micro	0.17	29	1.01 ± 0.46
	Nano	0.27	46	1.12 ± 0.19
	Pico	0.15	25	0.98 ± 0.15
	Total	0.59	---	1.05 ± 0.23
1026	Micro	1.96	59	0.48 ± 0.33
	Nano	0.99	30	1.55 ± 0.003
	Pico	0.39	12	2.00 ± 3.87
	Total	3.33	---	0.97 ± 0.30

Sta. CCB
(1290)

Date	P_m^B	I_k	ϵ_{PAR} (m^{-1})	Areal Prod ($mg\ C\ m^{-2}\ d^{-1}$)
2/24/09	3.2	21	0.7-1	1469/952
4/24/09	3.6	54	0.45	1523
2/10/10	1.8	23	0.66	424

Sta. 341

Date	P_m^B	I_k	ϵ_{PAR} (m^{-1})	Areal Prod ($mg\ C\ m^{-2}\ d^{-1}$)
2/24/08	3.4	29	1.32	230
2/17/09	2.1	9	0.68	313
2/27/09	3.2	14	0.68?	114

P_m : $g\ C\ g\ chl\ a^{-1}\ h^{-1}$

I_k : $\mu mol\ photons\ m^{-2}\ s^{-1}$

SUMMER RATES FOR COMPARISON

- Smith et al. 2005: 800-1200 $mg\ C\ m^{-2}\ d^{-1}$
- Fahnenstiel et al. 1995:
 - pre-zebra mussel: 738 $mg\ C\ m^{-2}\ d^{-1}$
 - post zebra mussel: 342 $mg\ C\ m^{-2}\ d^{-1}$

ARE THE DIATOMS ACTUALLY GROWING?

- Incubate samples with PDMPO
- Fluorescent compound incorporated along with newly deposited Si in diatoms frustules

(Leblanc and Hutchins, 2005 L&O:Methods)

- **Quantitative**: Incorporated at 2800:1 molar ration (Si:PDMPO)
- **Taxonomic**: Can see which cells are growing

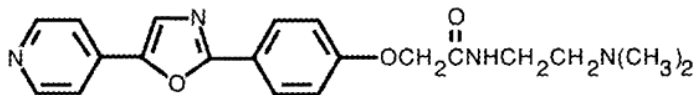
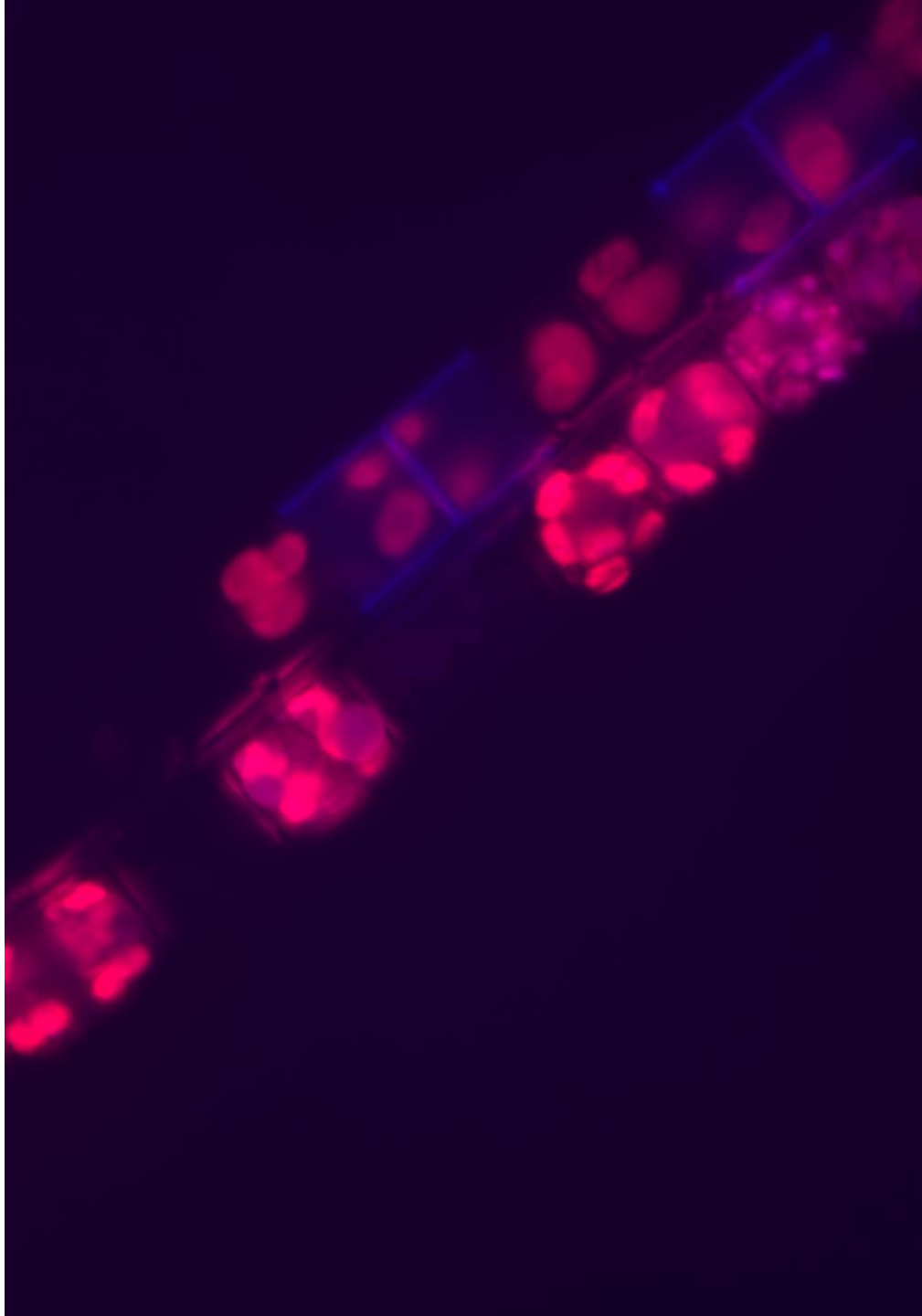


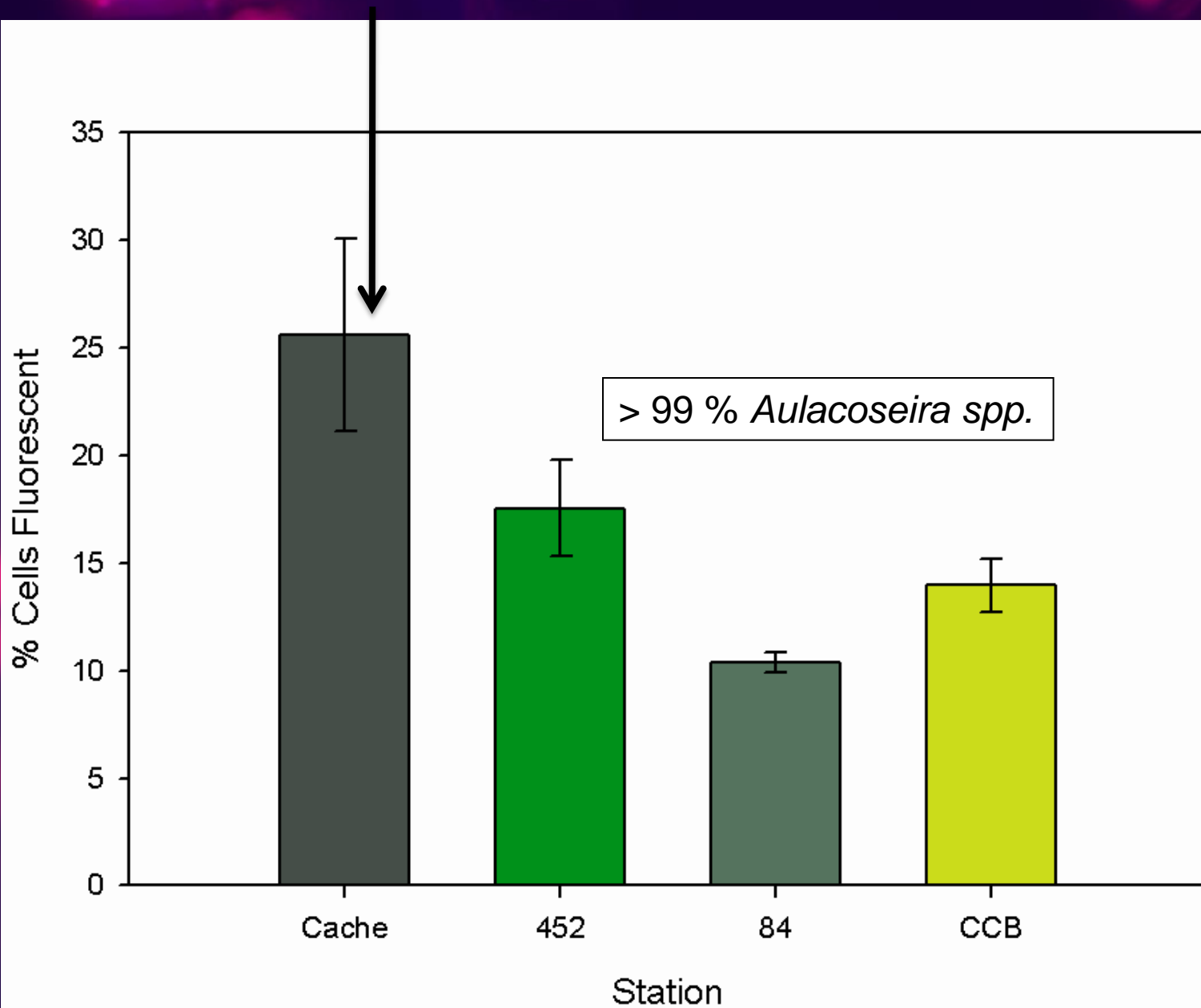
Fig. 1. Chemical structure of PDMPO [41].

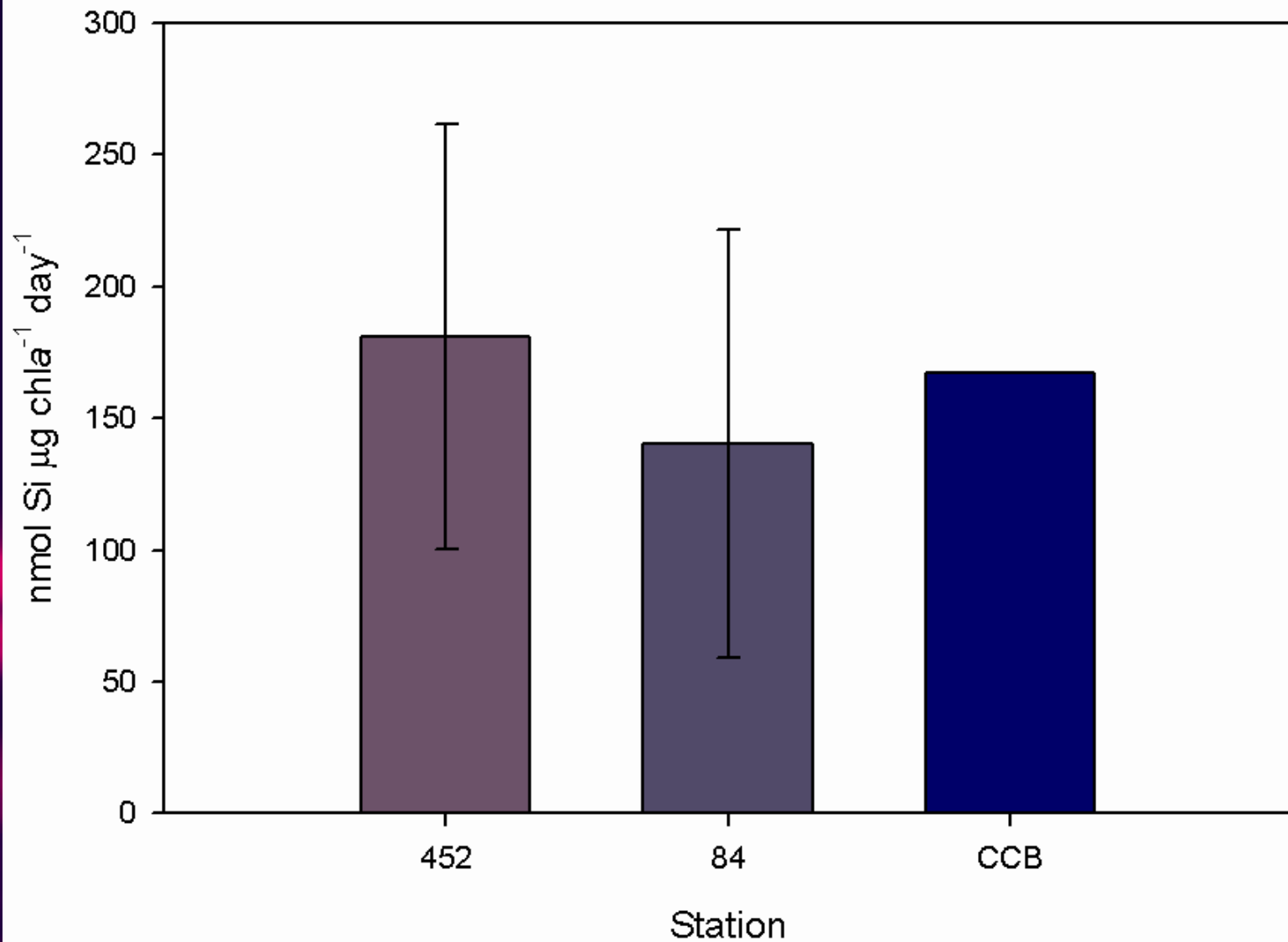


- Samples collected from water column
 - Incubated for 24 hours
 - Subsamples collected every 6 hours
-
- Microscopic determinations for taxonomy
 - PDMPO extracted and measured for rates



~3% *Cyclotella* spp. – all depositing Si





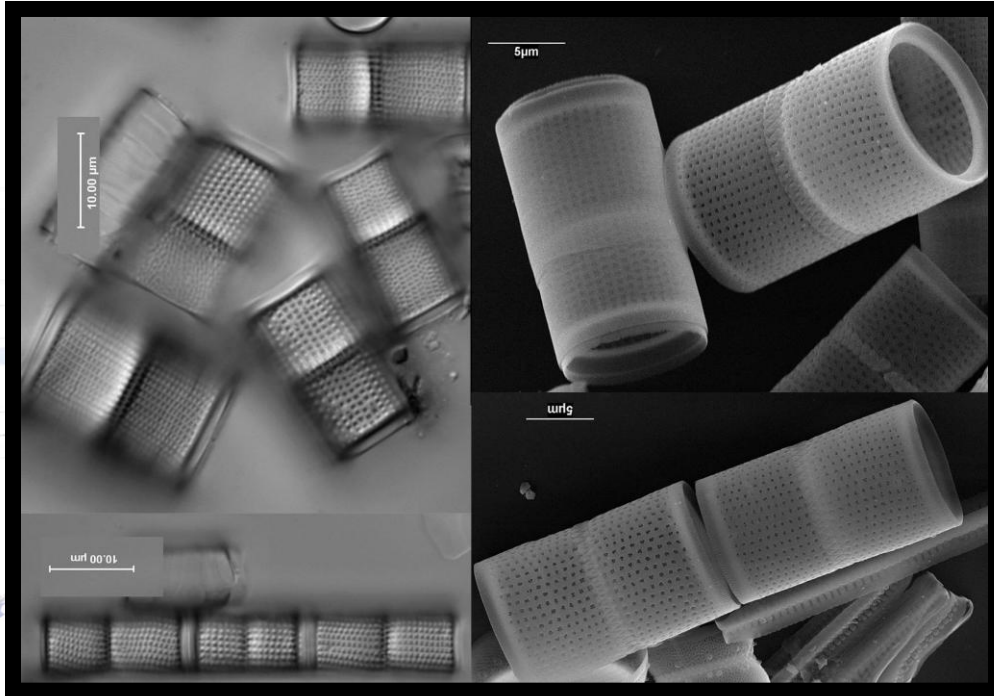
Aulacoseira islandica (O. Müller) Simonsen

Samples from Lake Erie 2007

Nutrient assimilation (molar)

Si:C = 0.03 – 0.04

Luxury C-fixation?

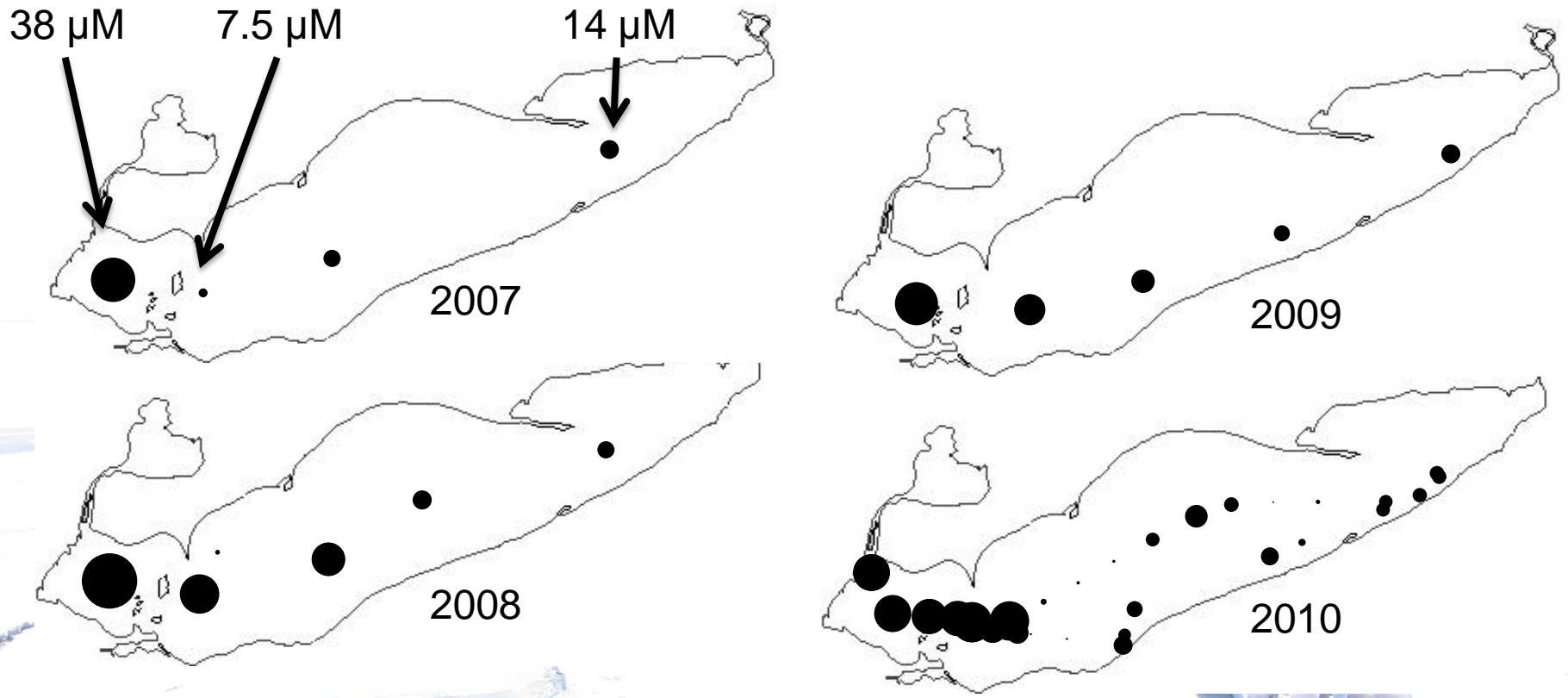


Photos by Furey and McKay, unpublished

**Brzezinski 1985
for 27 marine diatoms**

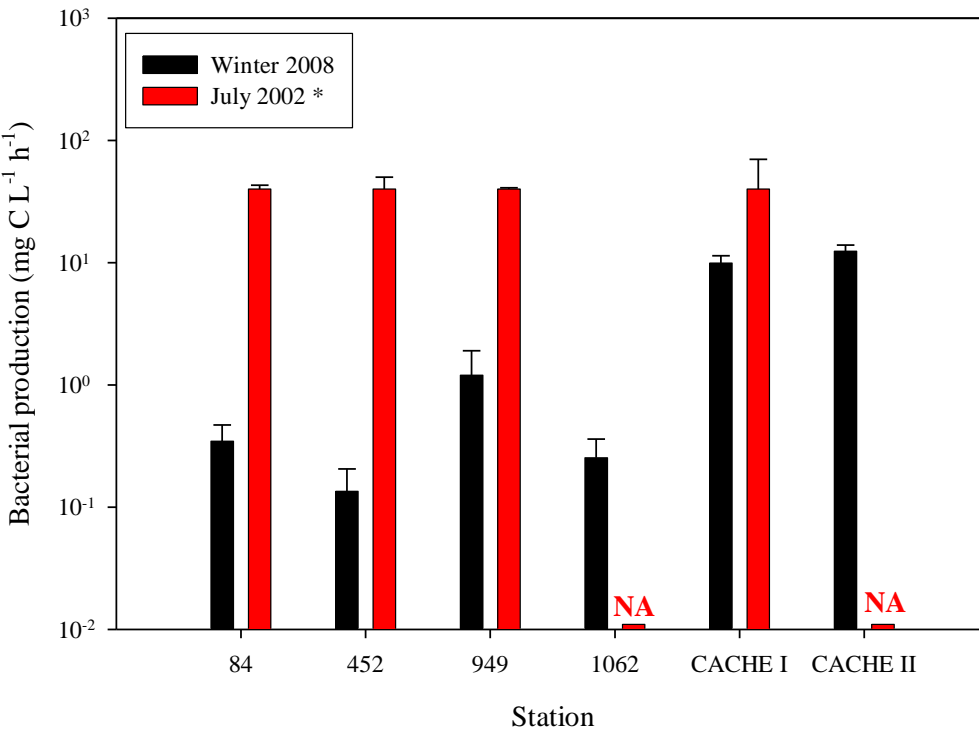
**Si:C = 0.13 ± 0.04
Range (0.04 – 0.36)**

Bubble plot of SiO₂ concentrations in January / February, 2007-2010



SiO₂ is markedly drawn down but never depleted

Surface bacterial production



Bacterial (2°) carbon production in the winter is 50 to 100 fold lower than in the summer.

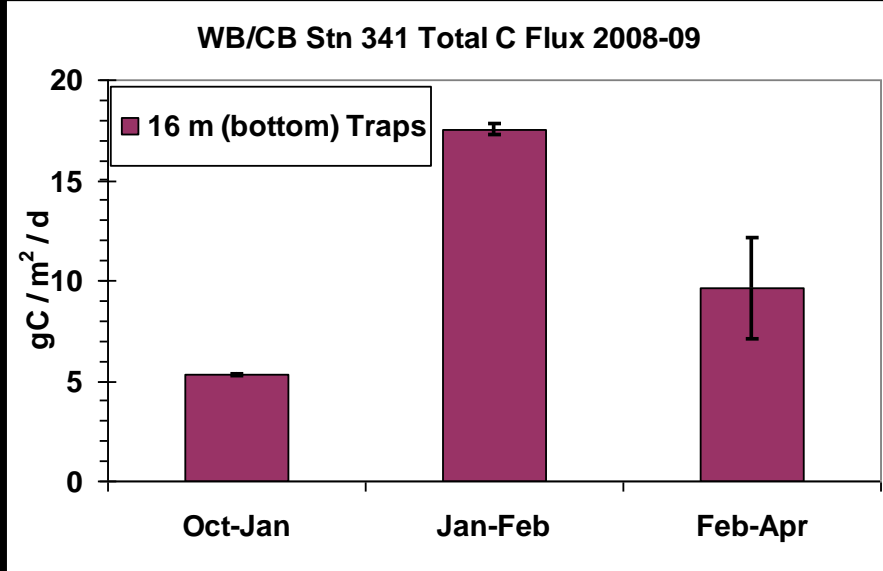
CACHE bacterial production is also lower than in summer months

Bacterial production is 0.011 % (n=14) of the average photosynthetic carbon production in winter months. CACHES (n = 2) 2° production is *ca* 0.27%.



Where does this winter biomass go?

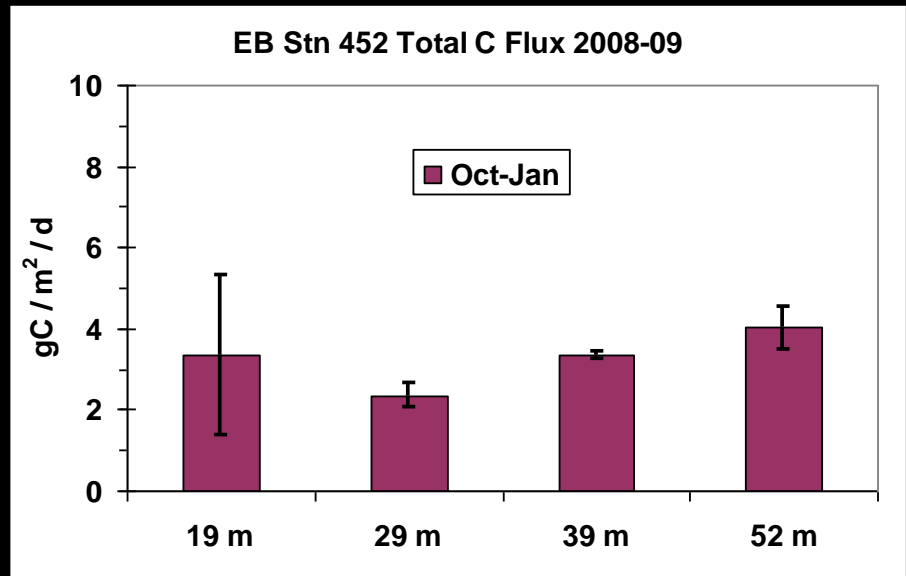
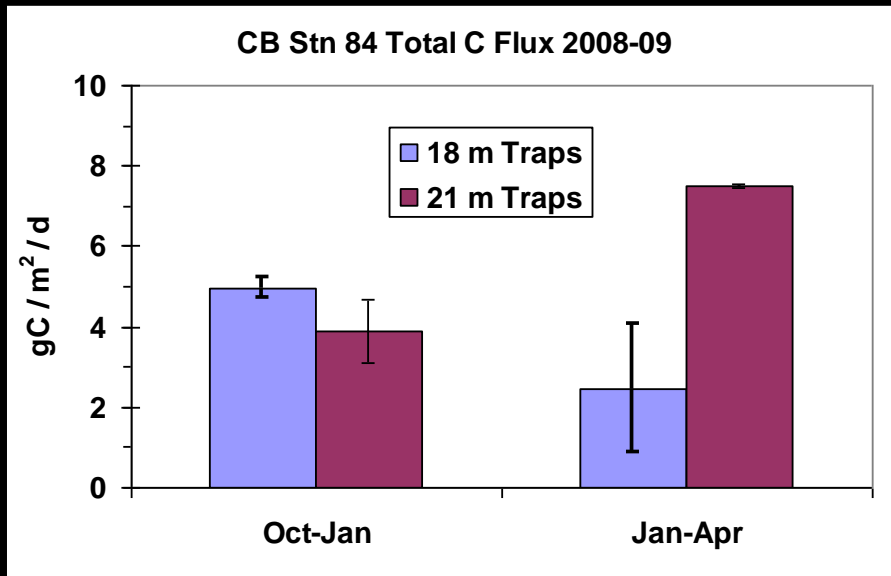
Total C Flux from Oct 2008 – Apr 2009



Stn 341 (note higher scale)
– evidence that winter C-flux is considerably higher than fall

Stn 84 - evidence for sequential settling of C, higher in winter

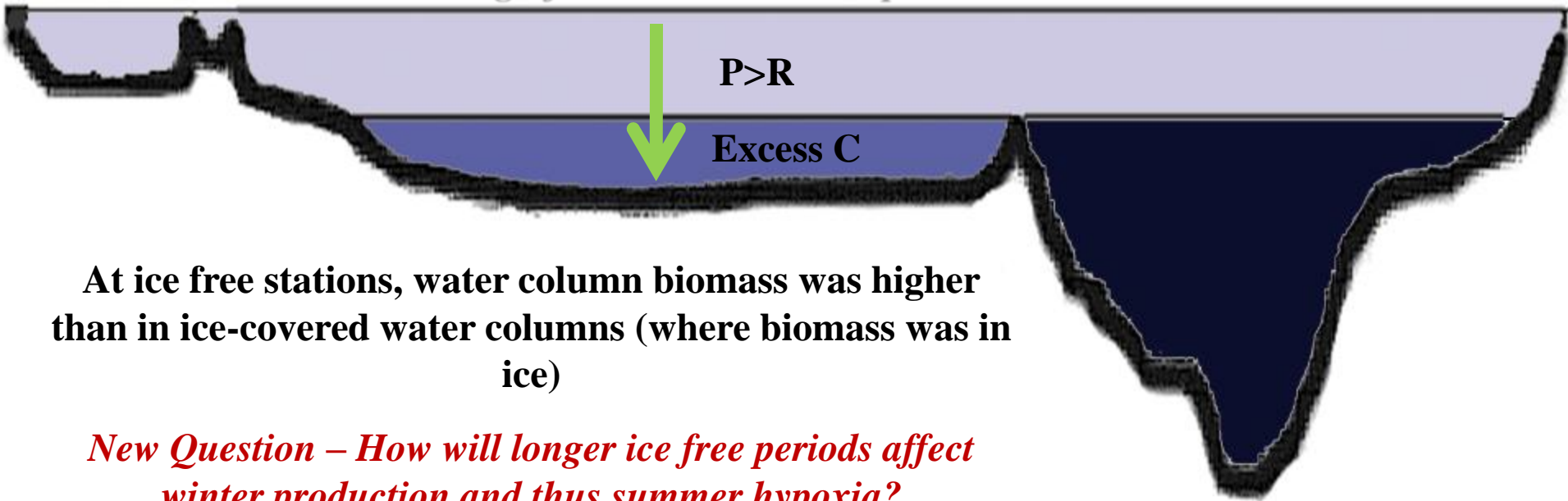
Stn 452 fall only - same range as other stations during fall



Summary

- Abundant photosynthetically “happy” diatoms dominate
- Significant wintertime carbon production and export in the water column
- Slow microbial decomposition rates

Significant water column production



At ice free stations, water column biomass was higher than in ice-covered water columns (where biomass was in ice)

New Question – How will longer ice free periods affect winter production and thus summer hypoxia?

