

## UKOA Sea Surface Consortium

# Initial results from the UKOA cruise to the Arctic, summer 2012.

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# Ocean Acidification Impacts on Sea Surface Biology, Biogeochemistry and Climate Consortium

## High-level objectives are:

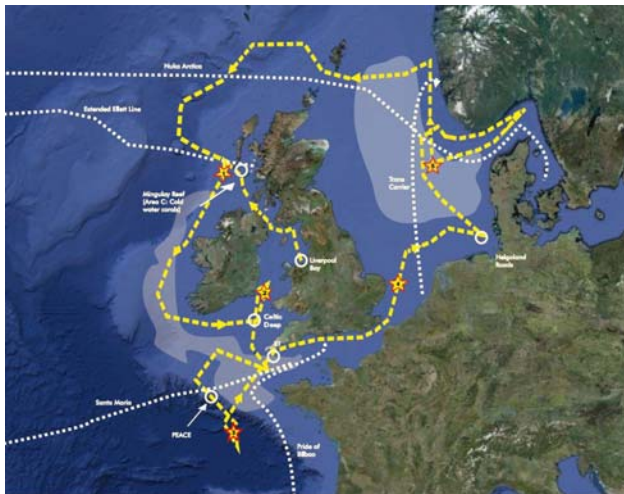
- To determine the impact of OA on **planktonic organisms** (in terms of physiological impacts, morphology, population abundances and community composition).
- To quantify the direct and indirect impacts of OA on **biogeochemical processes** affecting the ocean carbon cycle.
- To quantify the impacts of OA on the air-sea flux of **climate active gases** (DMS and N<sub>2</sub>O in particular).



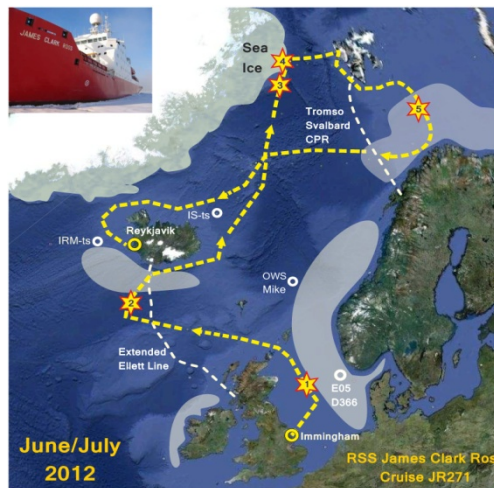
# Underlying Approach to Research

- **Comparative:** Observations of physical, chemical and biological ocean variables across a wide range of environmental conditions.
- **Experimental:** On-ship bioassays using natural plankton communities.

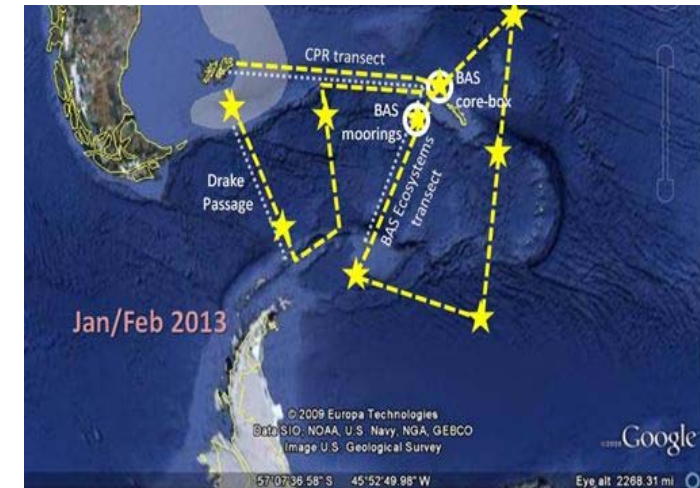
**European Seas  
June 2011**

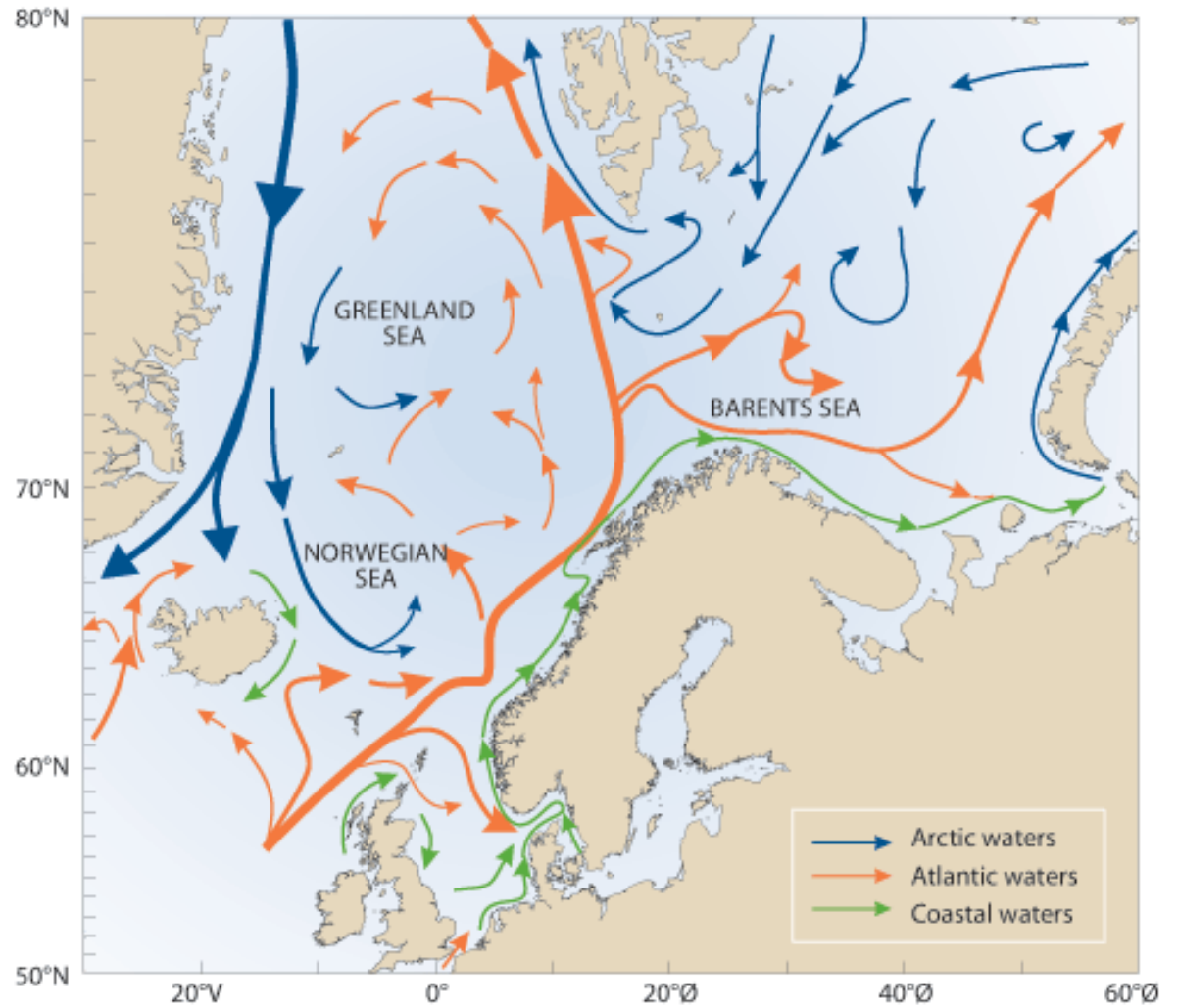
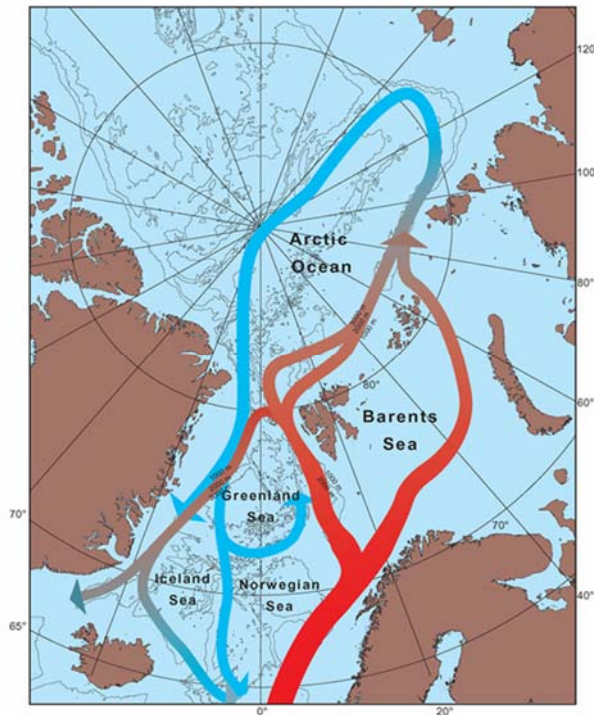


**Arctic Seas  
June/July 2012**



**Southern Ocean  
Jan/Feb 2013**





Source: North Sea Task Force, 1993. *North Sea Quality Status Report 1993*, Oslo and Paris Commissions, London, Olsen & Olsen, 132+vi pp.

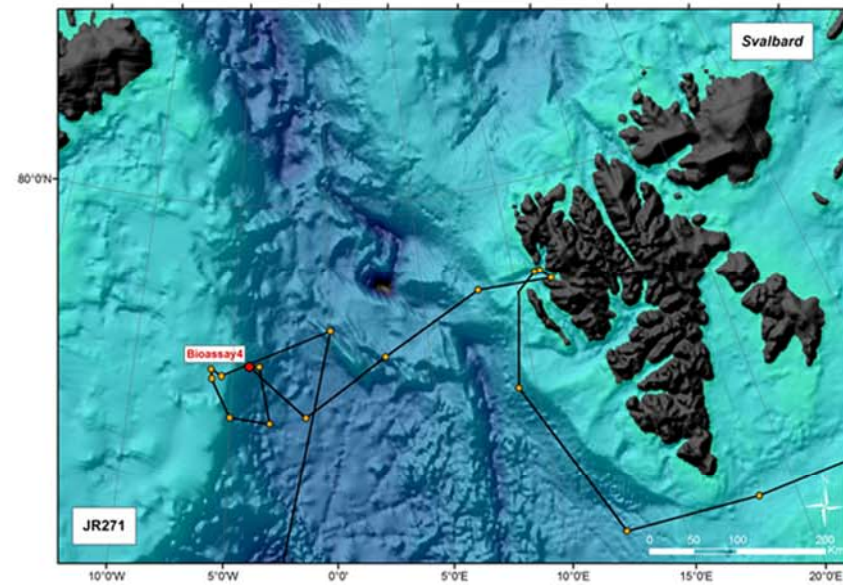
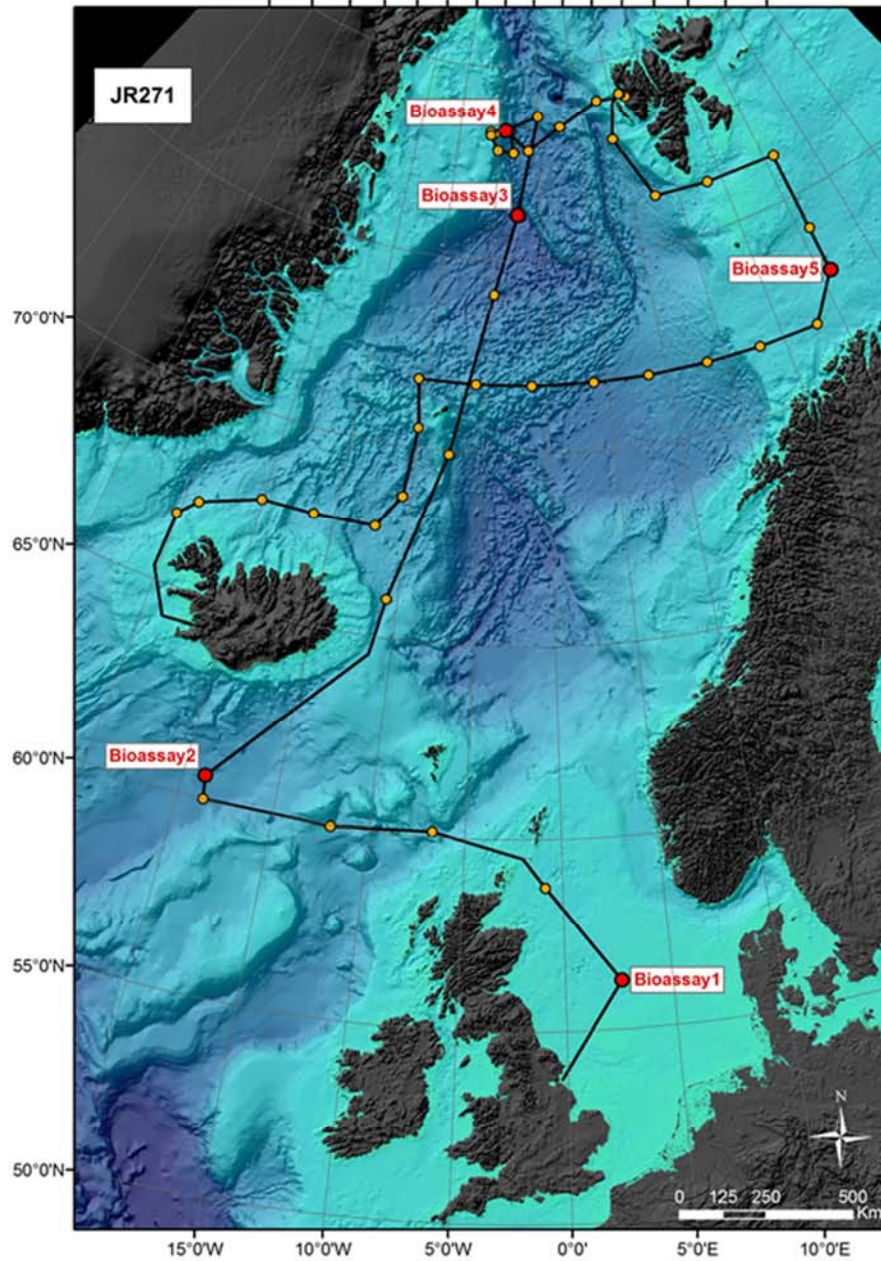


# Arctic Cruise Track

2 June to 2 July 2012



*RRS James Clark Ross*



# Comparative Observations

## Environmental Variables:

- Temperature
- Salinity
- Irradiance
- Carbonate chemistry
- Oxygen
- Macronutrients
- Micronutrients (Fe)

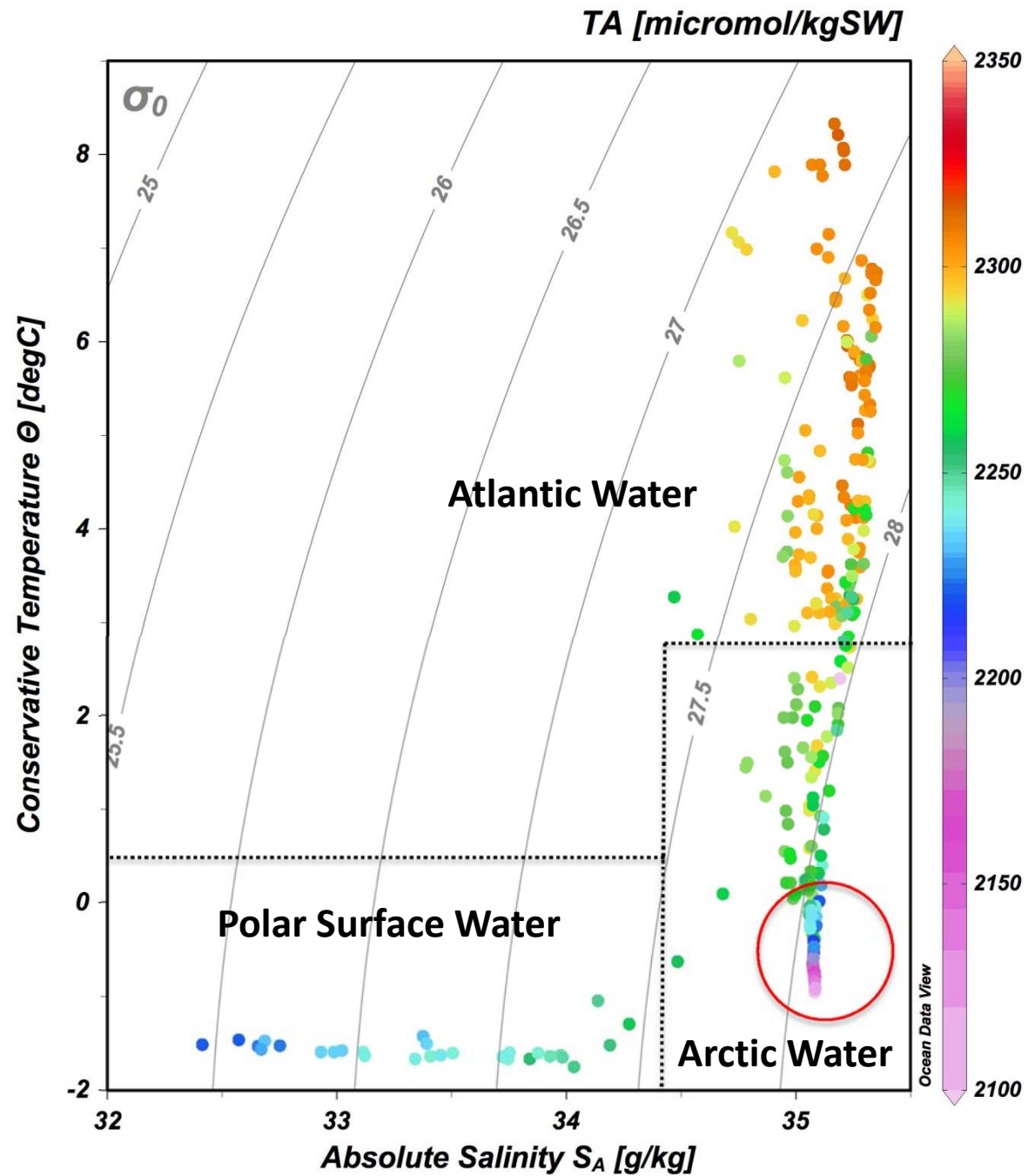
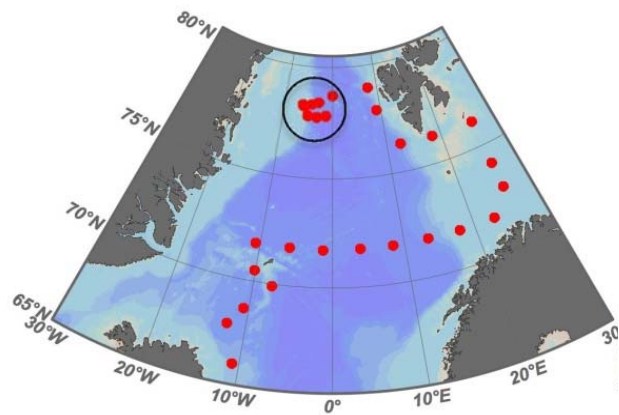


## Response Variables:

- Chlorophyll
- POC/N, DOC, TEP
- Photosynthesis
- Primary production
- Calcification
- Plankton Community
- Nitrification & N<sub>2</sub>O
- DMS and DMSP



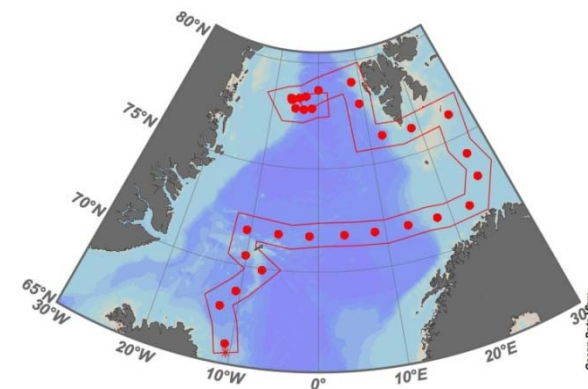
# Total Alkalinity



Data c/o E Tynan, M Ribas-Ribas and E Achterberg



# Distribution of $p\text{CO}_2$ ( $\mu\text{atm}$ )

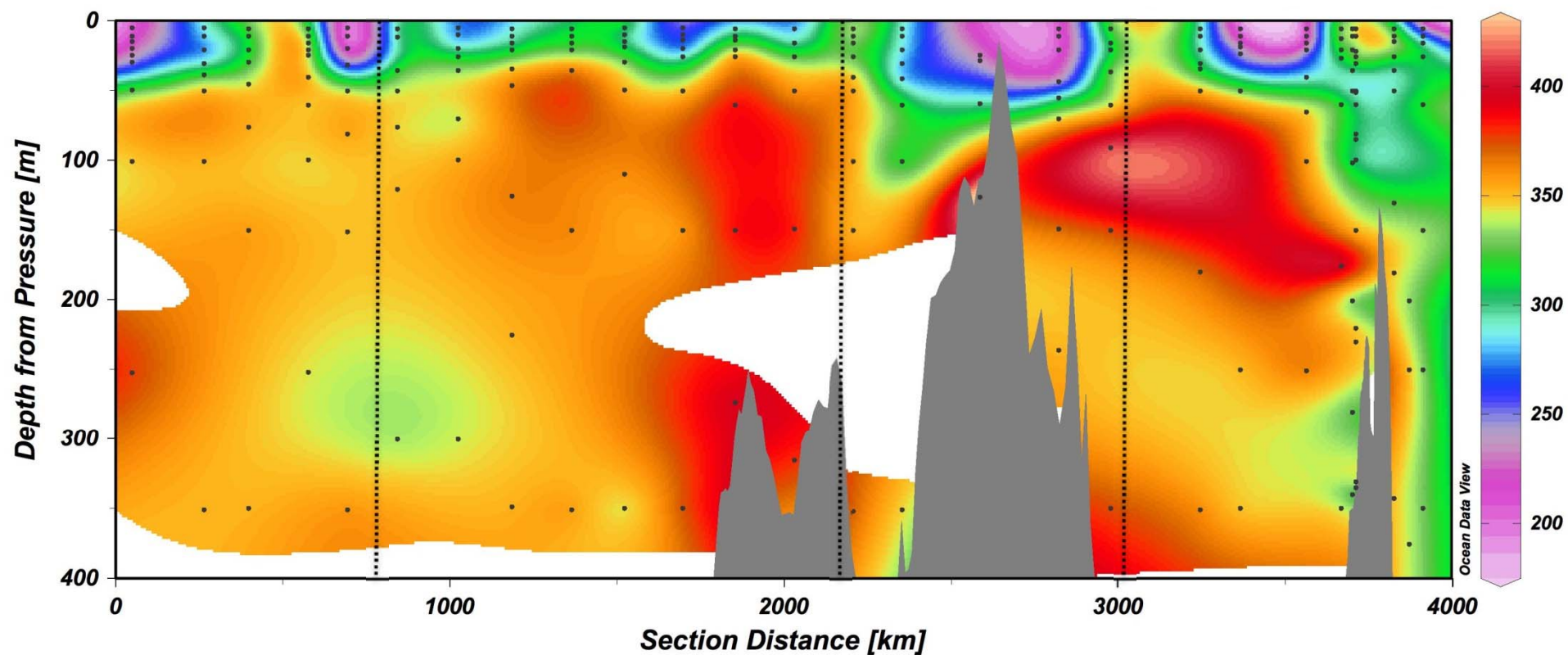


South of Jan Mayen

72°N Transect

Barents Sea

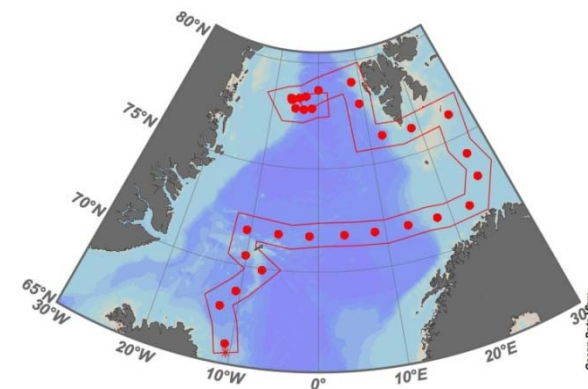
Fram Strait



Data c/o E Tynan, M Ribas-Ribas and E Achterberg



# Distribution of $\Omega_{ar}$

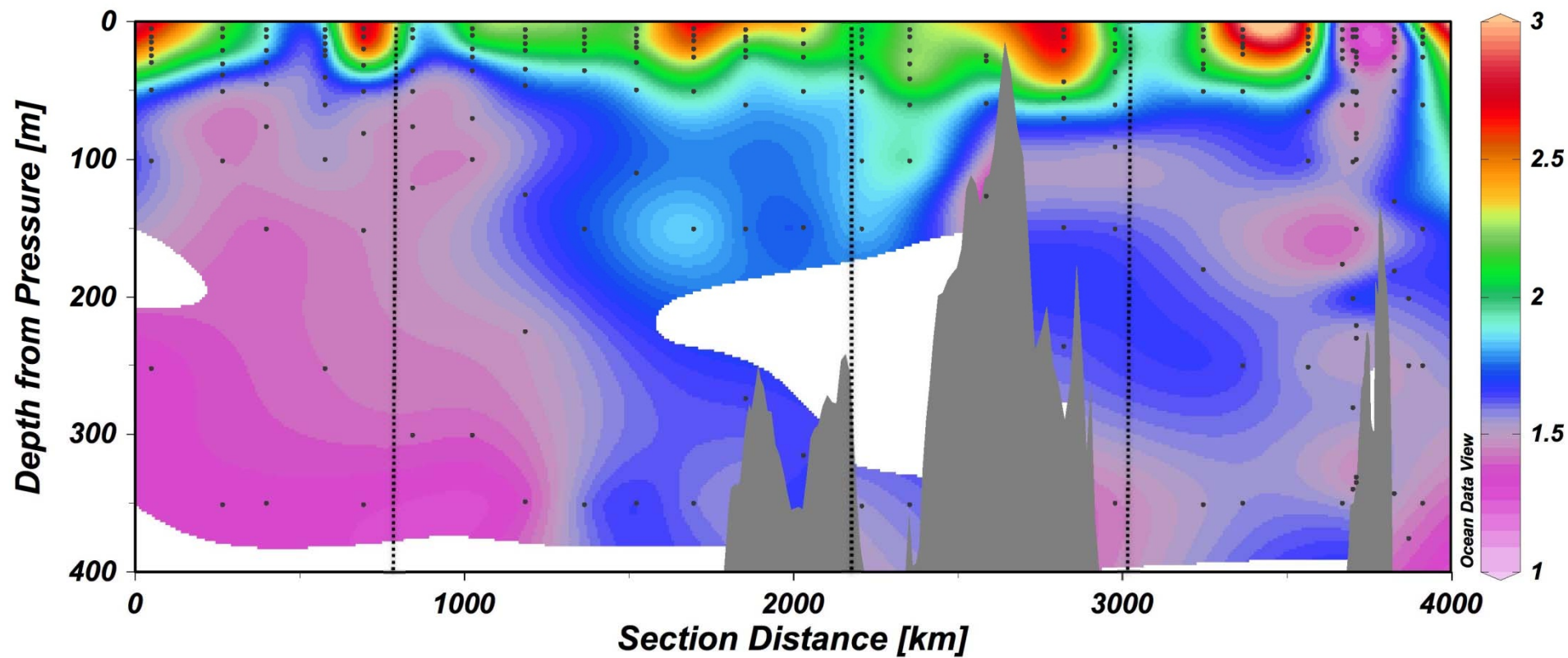


South of Jan Mayen

72°N Transect

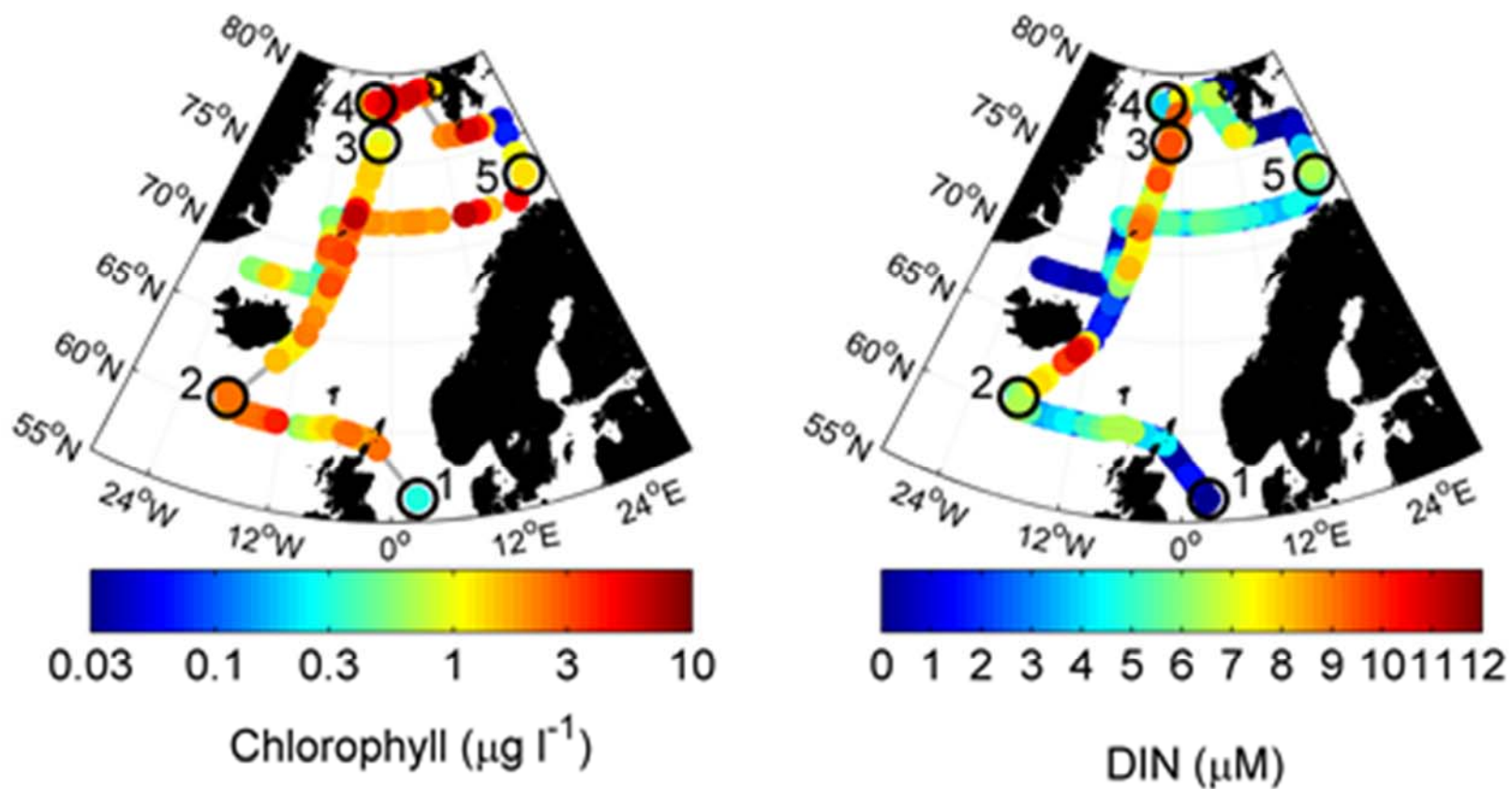
Barents Sea

Fram Strait



Data c/o E Tynan, M Ribas-Ribas and E Achterberg

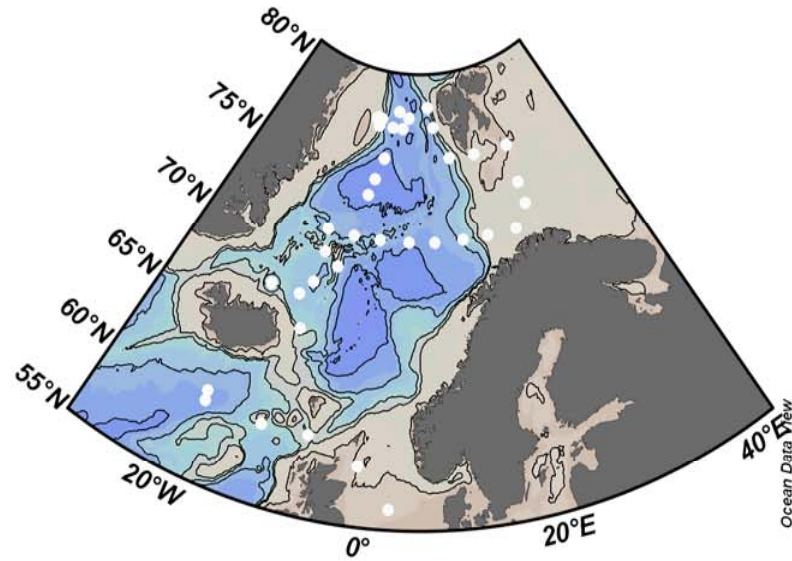
# Chlorophyll and Dissolved Inorganic Nitrogen



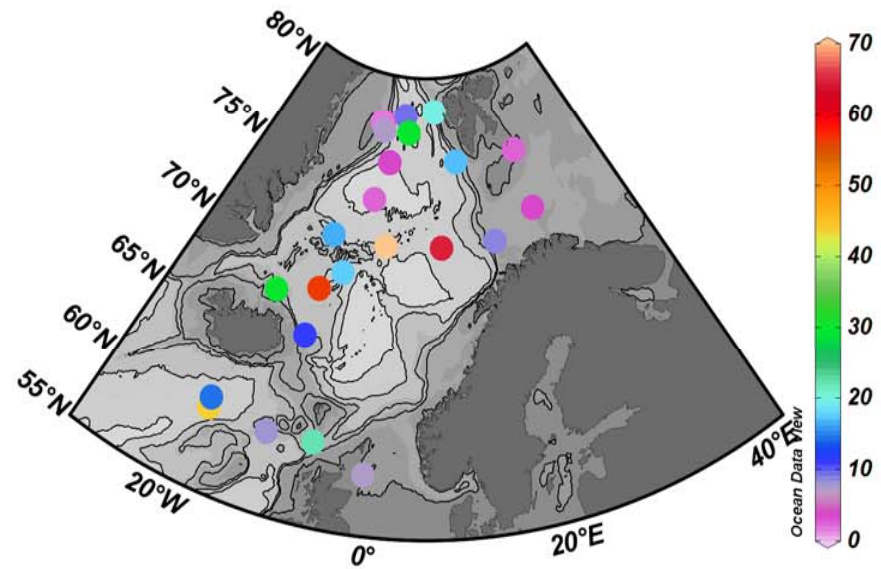
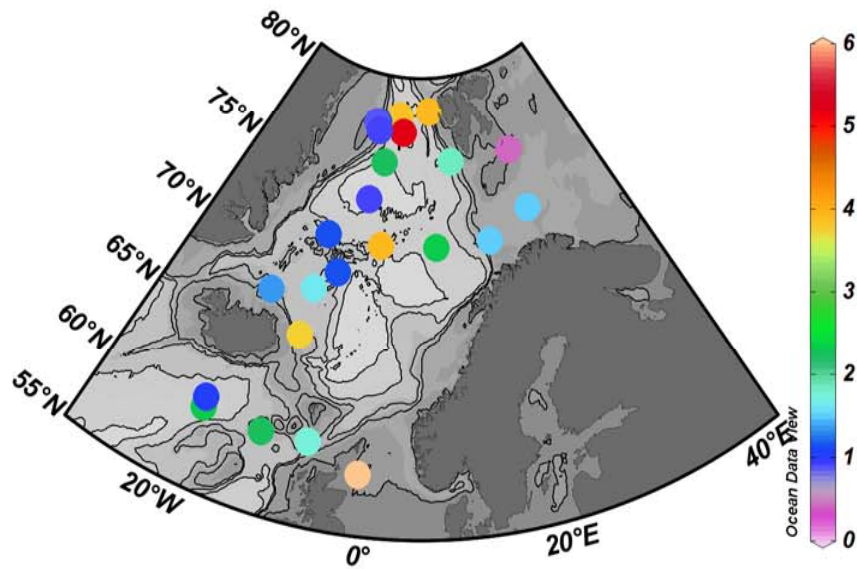
Data c/o S Richier, M Moore, A Poulton and M Esposito

# Primary Production and Calcite Production

Primary Production  
( $\text{mmol C m}^{-3} \text{d}^{-1}$ )



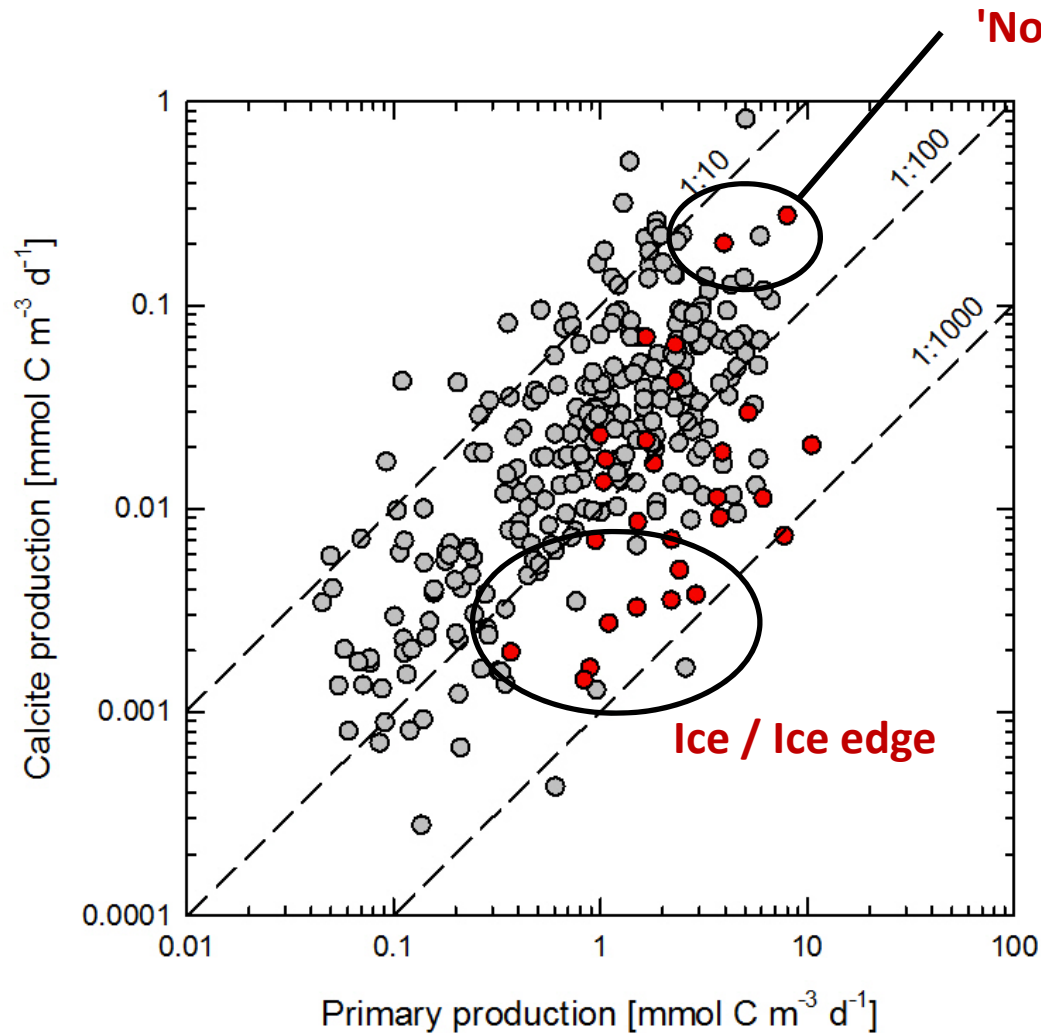
Calcite Production  
( $\mu\text{mol C m}^{-3} \text{d}^{-1}$ )



Data c/o A Poulton and C Daniels



# Calcite versus Primary Production

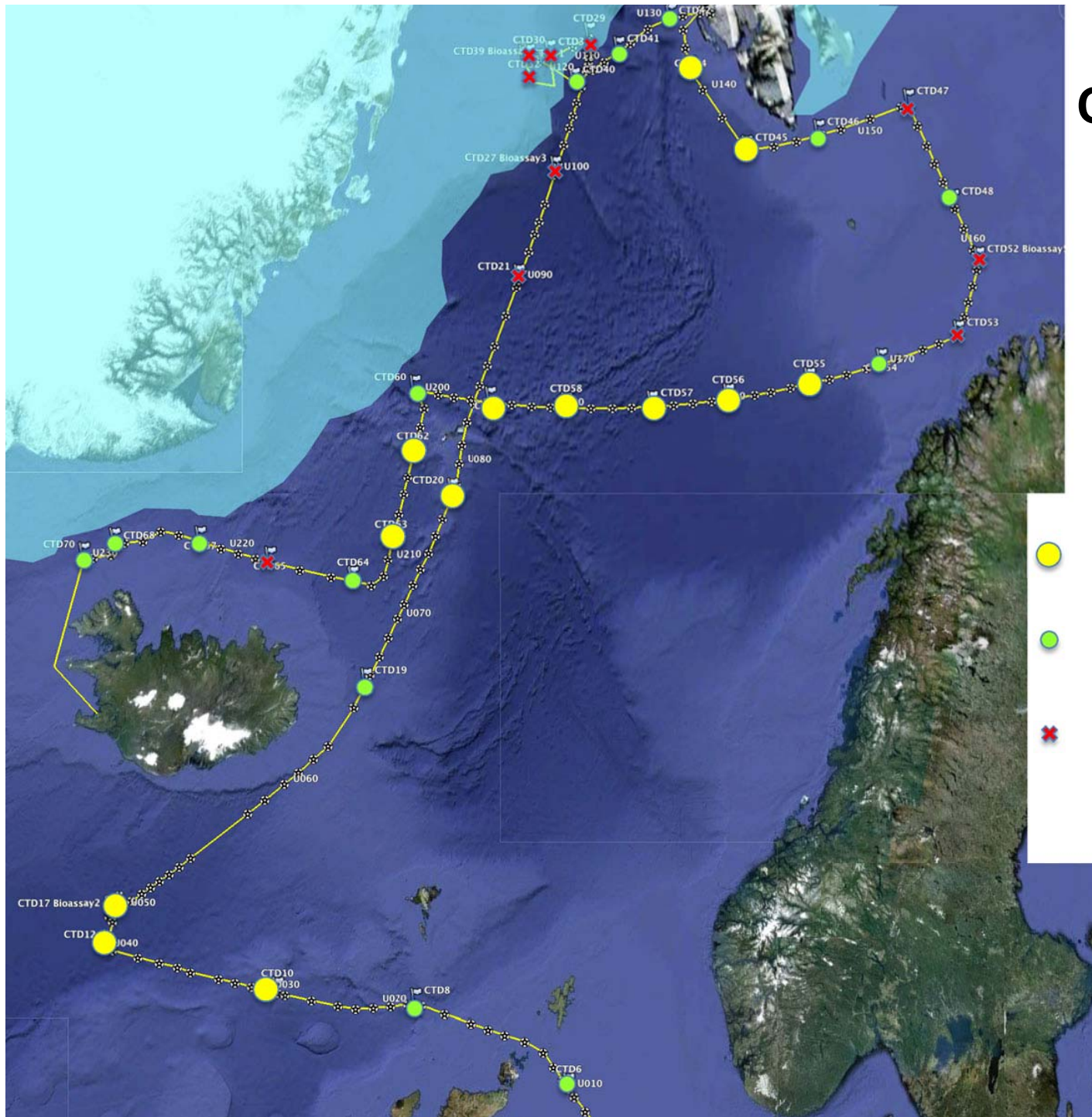


'Norwegian Sea'

- (Sub-)tropical Atlantic Ocean (AMT-14, 16 & 17)
- Iceland Basin 2007, 2010 (D321, D354)
- NW European Shelf waters 2011 (D366)
- Equatorial Pacific
- Patagonian Shelf
- Arctic and Drake Passage
- JR271 Arctic Ocean 2012

Earth System Science Data (ESSD),  
Collation of all discrete CP data  
(Global), Micro-Diffusion Technique  
~20 cruises, published / unpublished  
with Meta-data (analysis) 2013  
Poulton, Balch et al

# Coccolithophore Abundance



Data c/o J Young







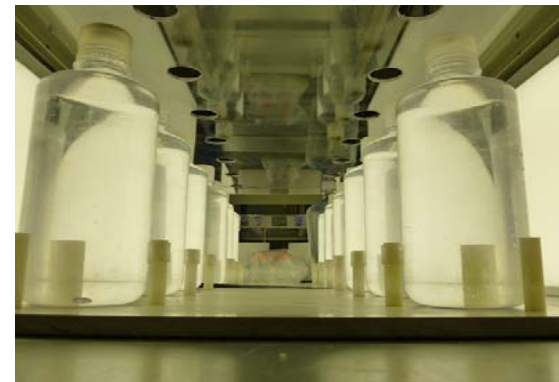
# Experimental Bioassays

Bioassay experiments, designed to evaluate the short-term response of multiple organisms and processes to artificial carbonate system manipulation.

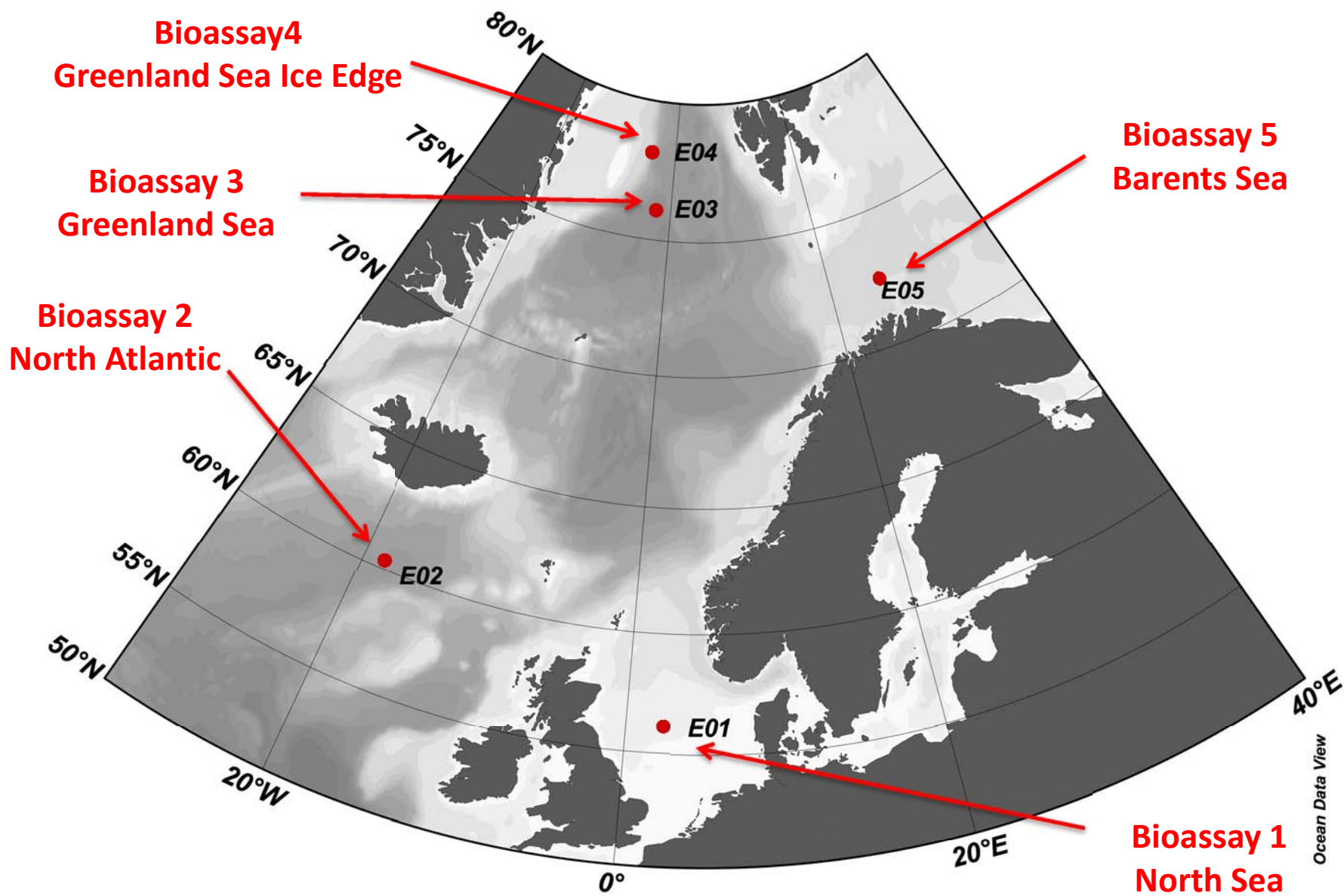
Bioassays were set up in different locations along the cruise track with different initial environmental conditions, reflecting both spatial variability within the study region and likely the temporal progression of the bloom.

## Experimental Design

- **Natural seawater samples collected:**  
3 replicate CTD Niskin Rosettes
- **pCO<sub>2</sub> manipulated (NaHCO<sub>3</sub> + HCL addition):**  
Ambient, 550 ppm, 750 ppm and 1000 ppm
- **Incubated on-ship at:**  
Light : 100  $\mu\text{E m}^{-2} \text{ s}^{-1}$  (14/8 L/D cycle)  
In situ temperature
- **Sampled at 3 time points:**  
0, 48h and 96h



# Results from Bioassay Experiments

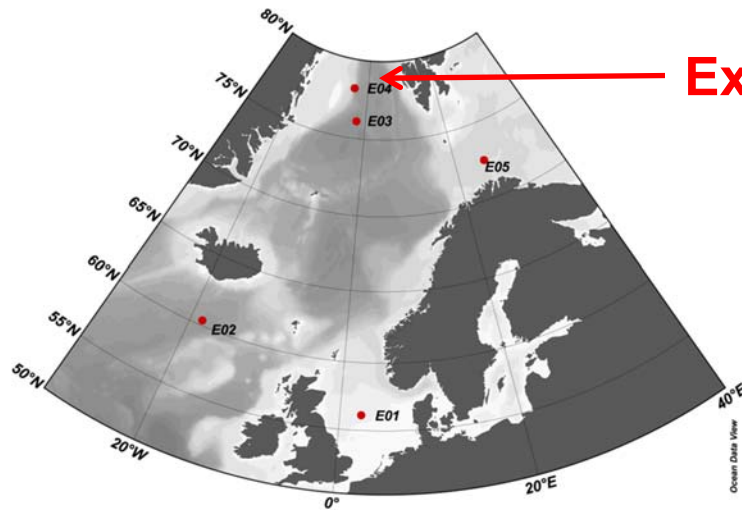


# Changes in Parameters in Control Samples after 48 & 96 hours

	Bioassay 1		Bioassay 2		Bioassay 3		Bioassay 4		Bioassay 5	
	North Sea		North Atlantic		Greenland Sea		Greenland Ice		Barents Sea	
	48 h	96 h	48 h	96 h	48 h	96 h	48 h	96 h	48 h	96 h
Nitrate	↓	↑	↓	↓	↓	↓	↓	↓	↓	↓
POP	↔	↑	↑	↑	↑	↑	↑	↑	↑	↑
Total Chlorophyll	↔	↑	↑	↑	↑	↑	↑	↑	↔	↑
>10 µm Chlorophyll	↔	↔	↔	↔	↔	↔	↑	↔	↓	↑
<10 µm Chlorophyll	↑	↓	↑	↑	↑	↓	↓	↓	↑	↑
PNAN Abundance	↑	↑	↔	↔	↓	↓	↔	↑	↑	↑
Bacterial Abundance	↑	↑	↓	↓	↓	↓	↔	↑	↓	↓
Bacterial Production	↔	↑	↓	↓	ND	↔	ND	↔	↔	↓
HNAN Abundance	↑	↔	↔	↔	↑	↑	↔	↑	↔	↑
DMS	↔	↑	↑	↑	↔	↑	↔	↔	↔	↑
DMSP	↓	↑	↑	↑	↓	↑	↑	↑	↔	↑
Nitrous Oxide	↑	↔	↑	↔	↑	↔	ND	ND	ND	ND



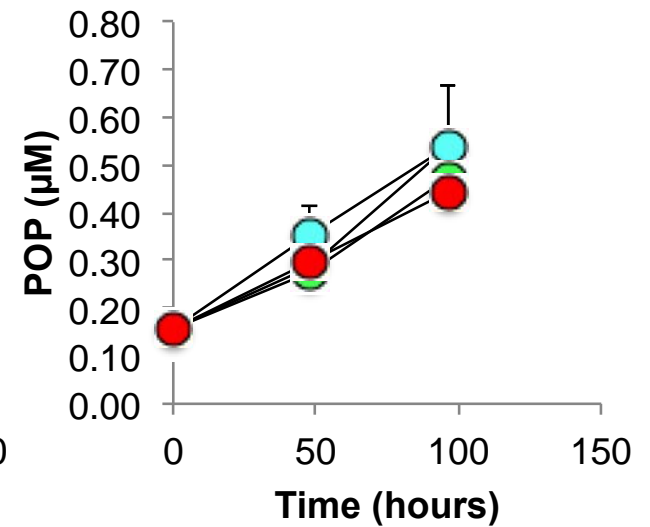
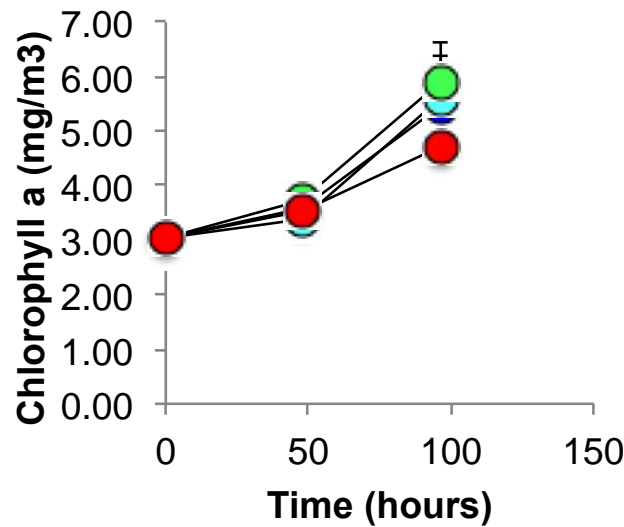
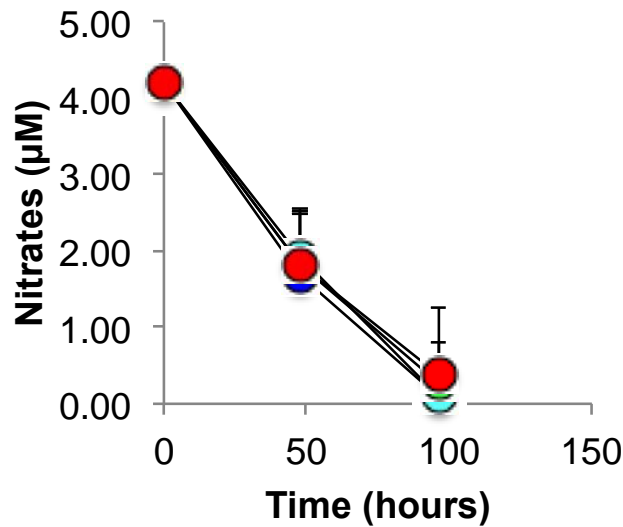
# Nitrate, Chlorophyll and POP Dynamics within Bioassays



**Example: Bioassay 4 Greenland Sea Ice Edge**

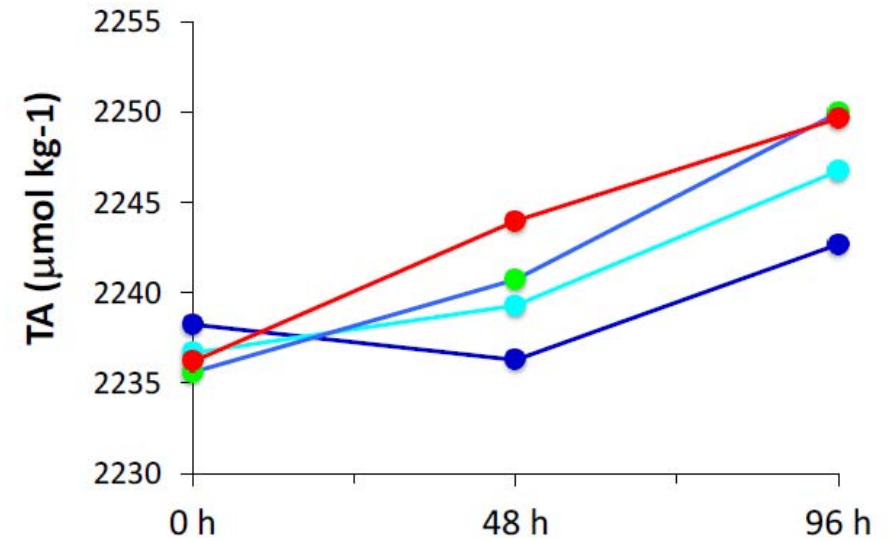
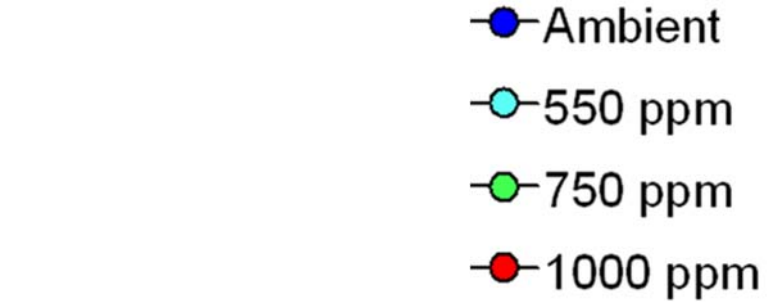
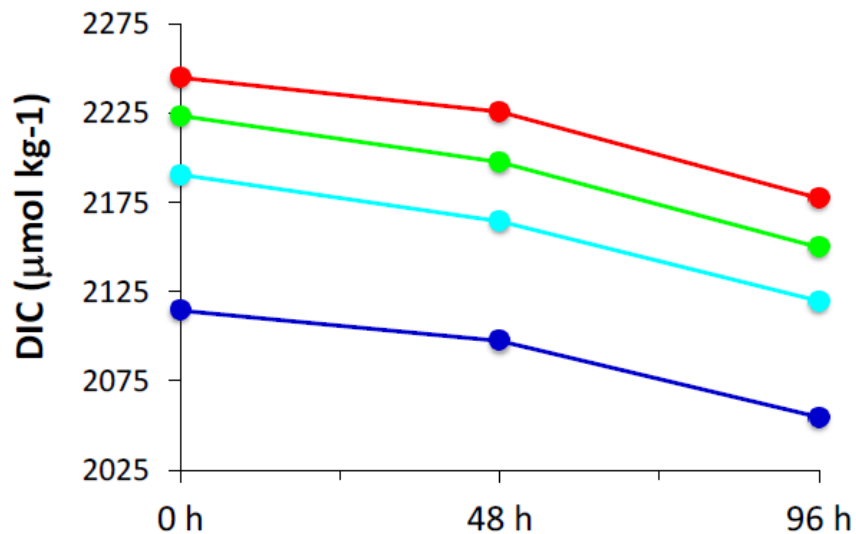
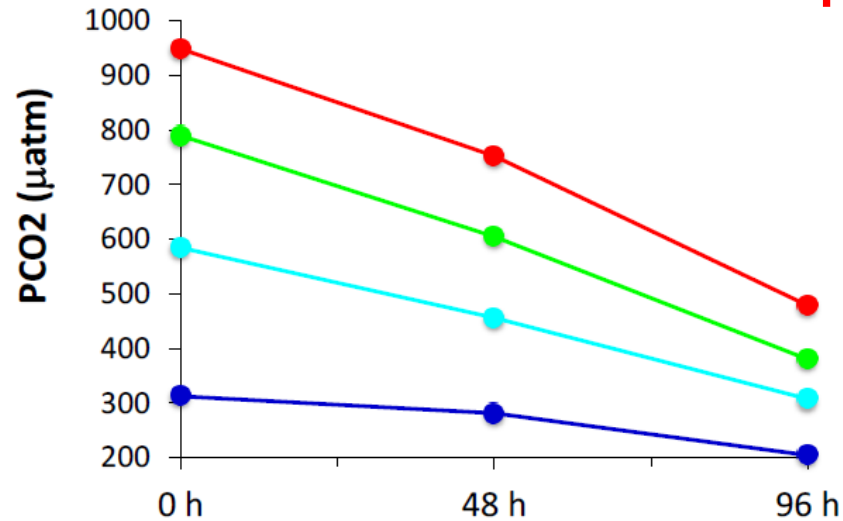
Temp (°C): -1.76  
 pH: 8.133  
 pCO<sub>2</sub> (ppm): 295.7  
 Ω<sub>calc</sub>: 2.38  
 Ω<sub>ar</sub>: 1.49

- Ambient
- 550 ppm
- 750 ppm
- 1000 ppm



# Carbonate Chemistry Dynamics within Bioassays

## Example: Bioassay 4 Greenland Sea Ice Edge



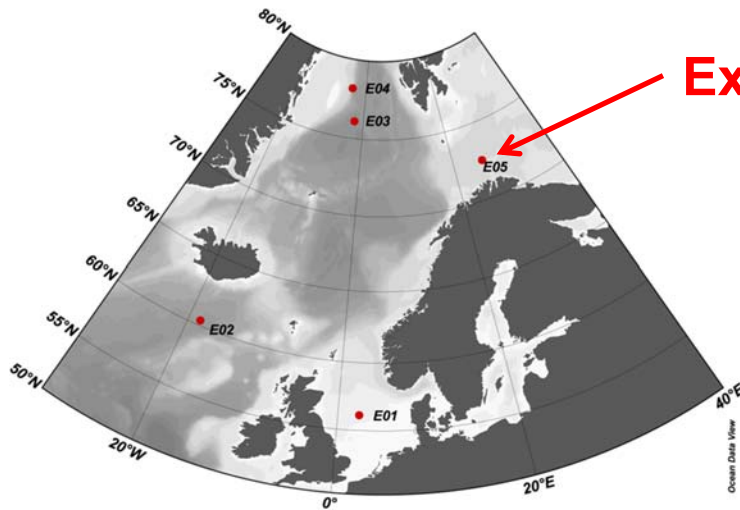
Data c/o E Tynan, M Ribas-Ribas and E Achterberg

# Response of Parameters to Acidification after 48 & 96 hours

	Bioassay 1		Bioassay 2		Bioassay 3		Bioassay 4		Bioassay 5	
	North Sea		North Atlantic		Greenland Sea		Greenland Ice		Barents Sea	
	48 h	96 h	48 h	96 h	48 h	96 h	48 h	96 h	48 h	96 h
Nitrate	-	-	-	-	-	-	-	-	-	-
POP	-	-	-	-	-	-	-	-	-	-
Total Chlorophyll	-	-	-	-	-	-	-	-	↑	↑
>10 µm Chlorophyll	-	-	-	-	-	↑	-	-	-	-
<10 µm Chlorophyll	↓	-	-	-	-	-	-	↑	-	-
PNAN Abundance	-	-	-	-	-	-	-	-	-	-
Bacterial Abundance	↑	-	-	-	-	-	-	-	-	-
Bacterial Production	↑	-	-	-	ND	-	ND	-	-	-
HNAN Abundance	-	-	-	-	-	-	-	-	-	-
DMS	-	-	-	-	-	-	-	-	-	-
DMSP	↑	-	-	-	-	-	-	-	↓	-
Nitrous Oxide	↓	-	↓	-	↓	-	ND	ND	ND	ND



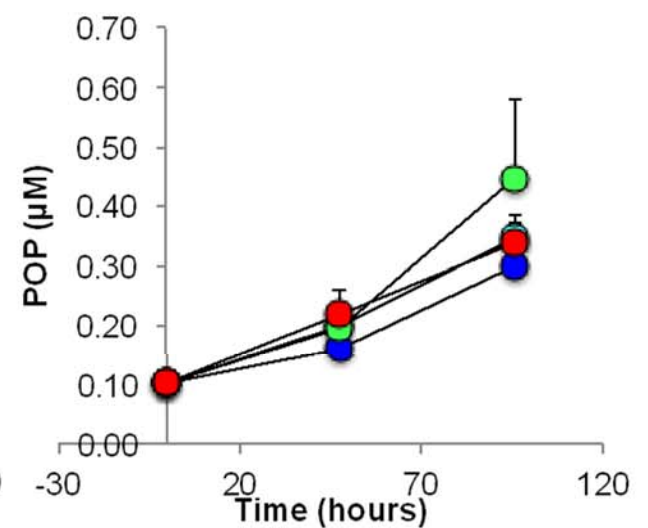
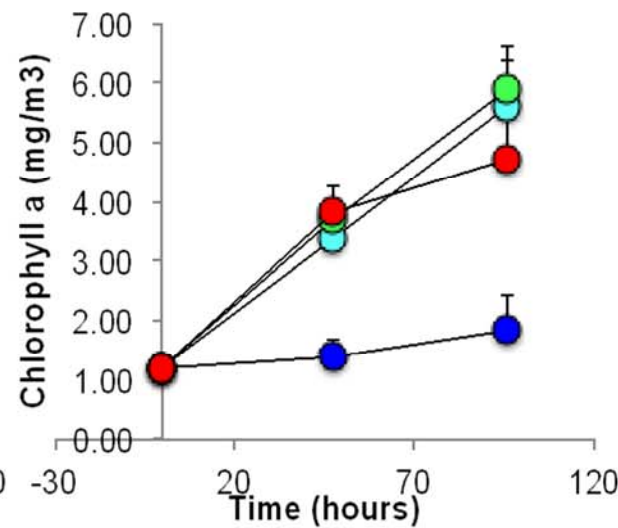
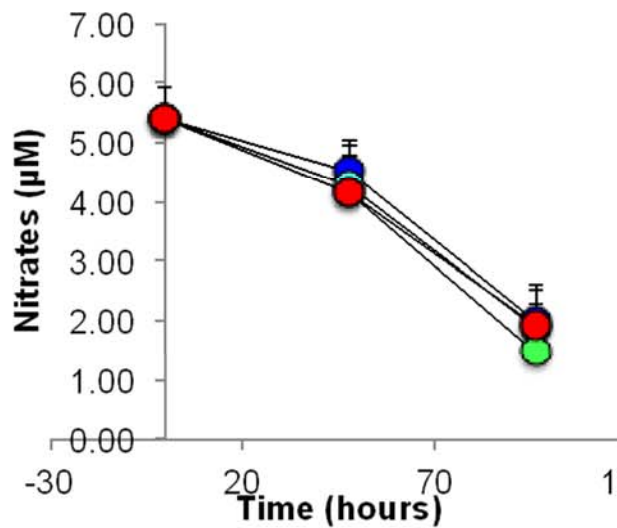
# Nitrate, Chlorophyll and POP Dynamics within Bioassays



## Example: Bioassay 5 Barents Sea

Temp (°C): 6.01  
 pH: 8.14  
 pCO<sub>2</sub> (ppm): 302.1  
 Ω<sub>calc</sub>: 3.48  
 Ω<sub>ar</sub>: 2.29

- Ambient
- 550 ppm
- 750 ppm
- 1000 ppm

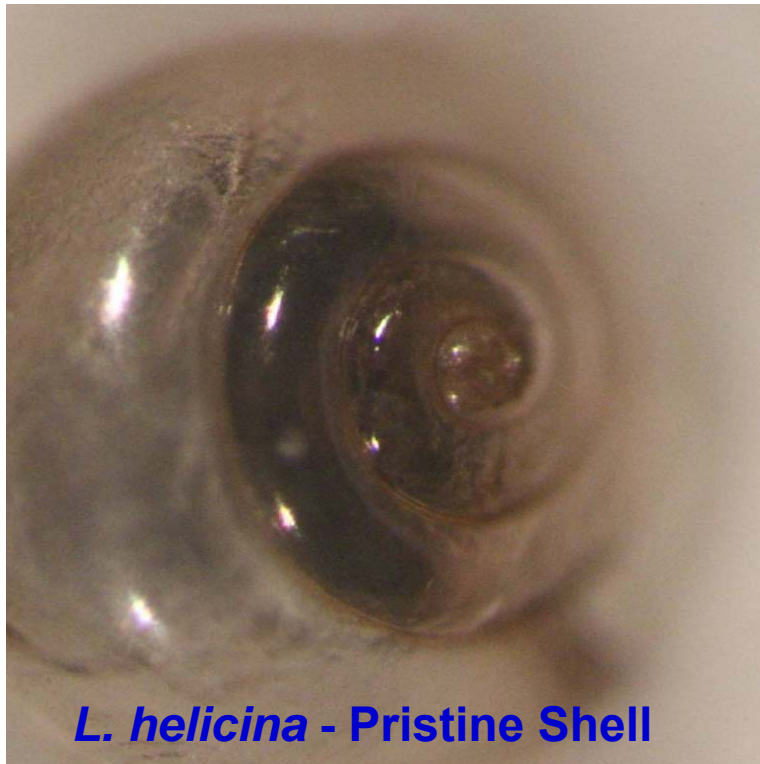


# Effect of Acidification on Pteropods

*Limacina helicina* collected from Kongsfjorden, Svalbard

Incubated for 8 days at Ambient, 550, 750, 1000 ppm pCO<sub>2</sub>

Examination of shell condition – damage and dissolution

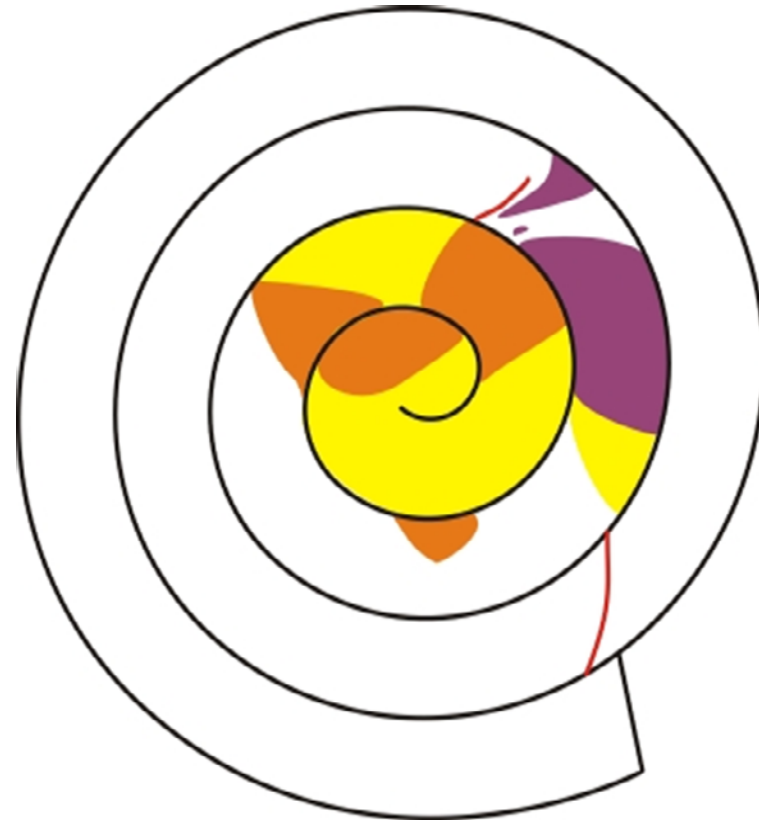
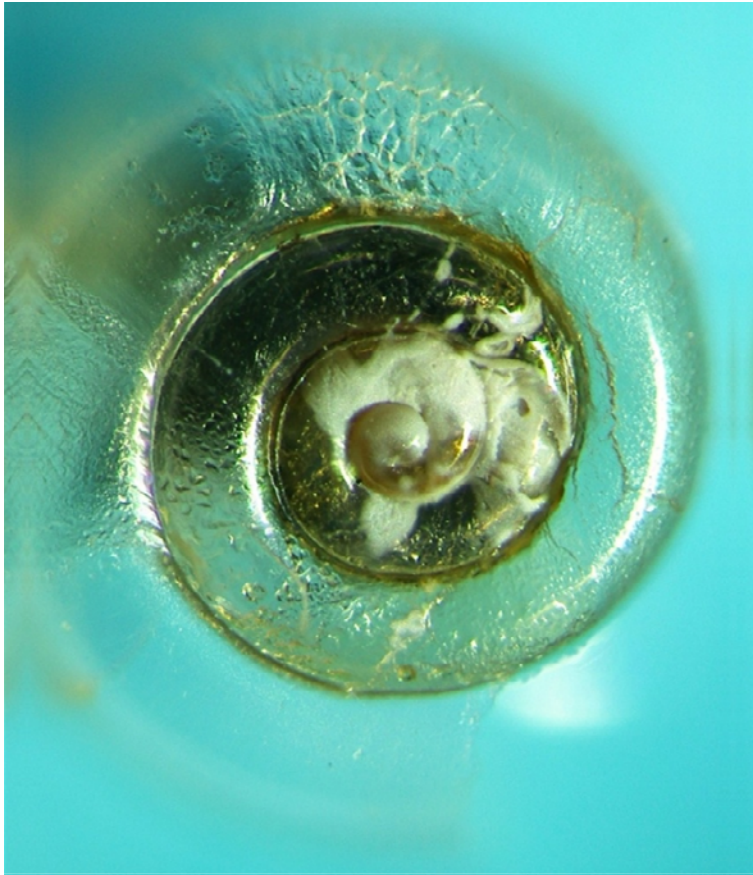


*L. helicina* - Pristine Shell



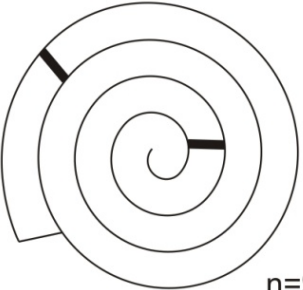
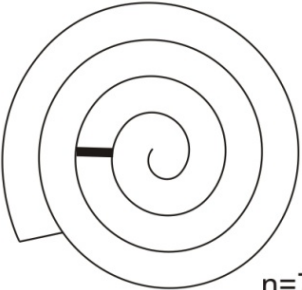
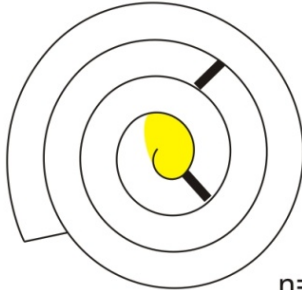
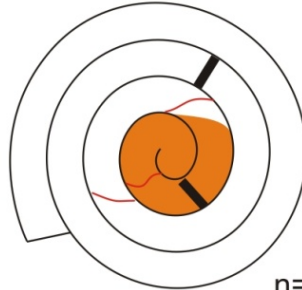
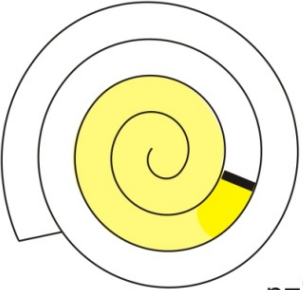
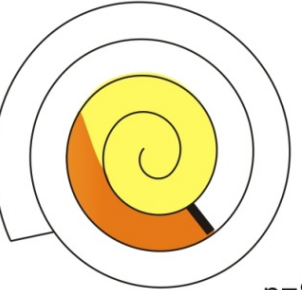
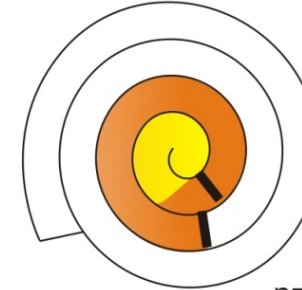
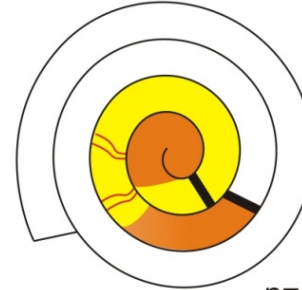
Damage to Shell Inner Whorls

# Imaging and Assessment of Shell Damage





## Effect of 8 Day Incubation on *L. helicina*

	Ambient	550 ppm	750 ppm	1000 ppm
ALIVE	 n=9	 n=7	 n=11	 n=13
DEAD	 n=7	 n=8	 n=10	 n=10

**Dissolution in living specimens was not observed below 750 ppm and localised to inner whorl(s)**

**Dissolution of dead specimens was present at ambient pCO<sub>2</sub> conditions and more uniformly distributed through shell**

# Conclusions

- **Wide range of observed variables measured with which to examine statistically the effects of carbonate chemistry.**
- **Biological response to incubation observed for most parameters measured to date in experimental bioassays.**
- **But no clear response to CO<sub>2</sub> treatment for most parameters measured to date in experimental bioassays**
- **Clear differential effect of CO<sub>2</sub> treatment on shell morphology of living and dead pteropods.**
- **These are preliminary results with much data still to be analysed.**

# Acknowledgements

Thanks to the captain, scientists and the crew of RSS *James Clark Ross* for assistance during the cruise.



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