DTU Aqua



Cruise report 09A12, R/V Dana, Hirtshals-Reykjavik, 27/8-2/9 2012 Oceanography at Sea DTU course 25501. Colin A. Stedmon, DTU Aqua cost@aqua.dtu.dk



Scientific Participants

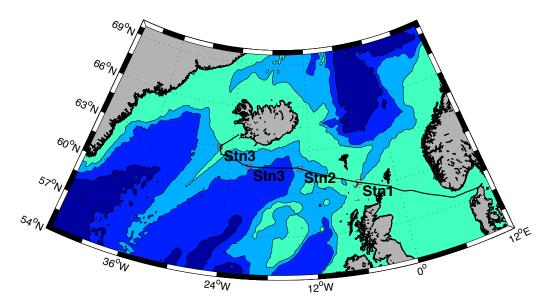
Colin Stedmon (Cruise leader, Course coordinator) Andre Visser (Teacher) Stiig Markager (Teacher) Katherine Richardson (Teacher) Tommy Nielsen (Technician) Nikolaj Thyssen Dam Lilli Gruwier Larsen. Karl-Søren Geertsen Thomas Bech-Thomassen Ciaran Joseph Murray Christina Søegren Stavroula A. Tsoukali Anette Christensen Søren Enghoff-Poulsen Arief Rullyanto Sanne Andrén Jakob Thyrring Kristiansen Mette Vodder Carstensen Haidi Cecilie Petersen Ida Margrethe Ringgaard Susan Guldberg Graungård Petersen Nikolaj Sørensen Nanna Finne Jensen

Cruise Objective

This cruise was the focus of a university course (5 ECTS) in oceanographic research techniques on board R/V Dana, sponsored by the Danish Centre for Marine Research. The course was designed to give students sea-going experience and a practical introduction to ocean sampling. The learning objectives of the course were:

- Identify common pitfalls and necessities with organizing oceanographic sampling
- Use a conductivity-temperature-depth (CTD) probe for measuring the physical properties of seawater
- Conduct measurements of underwater light penetration and assess the contribution of different water constituents to light attenuation.
- Quantify phytoplankton biomass distribution using chlorophyll a measurements and assess phytoplankton productivity.
- Conduct measurements of chemical properties of seawater (concentrations of oxygen and selected nutrients)
- Use ODV for processing and presenting oceanographic data.
- Integrate and interpret the oceanographic data collected.
- Relate the data collected during the cruise to regional oceanographic phenomena.

Figure 1: The route sailed and the location of the stations visited. At each station CTD (+light) profiles and water samples were collected.



Nature of work carried out on board.

The course was designed to have four oceanographic sampling themes: biological; physical; chemical and optical. The students were split into four groups and rotated between each theme each day. Biological oceanographic theme covered sampling



for phytoplankton and measuring chlorophyll concentrations using a spectrophotometer and carrying our Winkler oxygen measurements. Physical measurements included operating the CTD, coordinating water sampling between the groups, communicating with the bridge, and sampling for salinity calibrations. The chemical oceanography components covered sampling and analysis of pH, Alkalinity, phosphate and silicate concentrations, and the optical theme covered measurements of the attenuation of broadband light (PAR) in the surface waters and comparing these with the optical properties of particles and dissolved organic matter measured from water samples.

Cruise log

27th August,

All arrived in Hirtshals and boarded Dana. Equipment set up and secured in the laboratories. Sailed from Hirtshals around 14:00 (UTC) in good weather. Course started with a brief introduction and outline by Colin. After dinner, Andy gave a talk introducing some of the fundamentals of ocean physics. At around midnight the sea state began roughen and winds picked up.

28th August,

The main plan for the day was lectures while we steamed across the North Sea towards the first sampling station south of the Faroe Islands. There were high winds and 5 m swells making for challenging teaching conditions. During the day the conditions improved. During the morning introductory lectures on aspects of biological, chemical and optical oceanography where given by Katherine, Colin and Stiig, respectively. After lunch an introduction to the oceanography software Ocean Data View was given and the students carried out some exercises with the software.

29th August,

Swells picked up again a little in the night but by morning returned to relatively calm and overcast conditions. The morning was used to prepare for the first sampling in groups. We arrived at Station 1 in the Faroe-Scotland Channel at about 13:30 and started with a shallow CTD cast, collecting water for phytoplankton abundance and primary production, and bio optics measurements. Then carried out a deep CTD cast down to 1100 m, collecting water for water chemistry. Students worked on the samples and data collected while the ship then continued to steam west towards the Iceland Basin. It was a late night in the lab for some and a long night processing CTD data for others.

30th August,

Sea state fine, only a gentle swell, and the clouds where clearing. Arrived at Station 2 in the Iceland Basin at around 9 am. Carried out two casts again, a deep and a shallow. Students worked on their samples and data during the rest of the day. After dinner two Ph.D. students, Areif (DTU) and Ciaran (AAU) gave talks presenting their PhD projects.

31st August,

Arrived early in the morning at the third station south of Iceland. First cast with the CTD started at 6 am down to approximately 1000 meters. Followed by a shallow cast for phytoplankton samples and light measurements. Students busy the rest of the day in the lab although we were finished earlier due to the early start. Before dinner Nikolaj Sørensen (KU) gave a presentation of his PhD and after dinner Stavroula (DTU) presented her project. A whale was spotted in the distance at about 8 o'clock in the evening and everyone was up in the bridge on the look out. It did not come too close and it was difficult to sea its body or tail as there was quite a swell. In the evening the weather turned bad again with high winds and swell. A gentle reminder for the students that anything not strapped down or correctly stowed away goes flying!

1st September

Arrived on station southwest of Iceland at 5:30 in the morning. Same sampling plan carried out as the day before. In the afternoon, whilst steaming to Reykjavik, the students were then reorganized into the groups they will be working up data in, and began discussing with the teachers how to present and interpret the data and plans for eventually making a poster. After dinner, Stiig gave a short talk that he was preparing for the Danish parliament on the status of nitrogen loadings to Danish coastal waters. After the talk we arrived in Reykjavik. The students continued in their groups working on the data collected from the cruise. Andy Visser disembarked.

2nd September

After breakfast, a short meeting and group picture, the students disembarked, tired but happy. The teaching cruise appeared to have been a success, with the students grasping the realities of sampling at sea and obtaining hands on experience with a range of techniques.

Findings.

To document the findings of the work the four posters created by students are in the appendix of this report.

Concluding remarks.

The course was a great success. The crew of Dana, the teachers and the students all enjoying the experience. I would like to take the opportunity to thank all involved and especially the Danish Centre for Marine Research for providing the funds to support the course. The teaching on board worked well and I highly recommend that this is repeated in the future. Dana and its crew provide an inspirational teaching and learning environment.

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Biological Oceanography at Sea

The aim of this cruise was to learn to organize biological oceanography data sampling. A conductivity-temperature-depth (CTD) probe measured physical, chemical, optic and biological properties of North Atlantic water and we used the sampling results to quantify phytoplankton biomass distribution and productivity. Some CTD-plots are presented in the software program OceanDataView (ODV). We focus on the different factors influencing phytoplankton productivity and distribution and in addition discuss the changing conditions in the world's oceans.

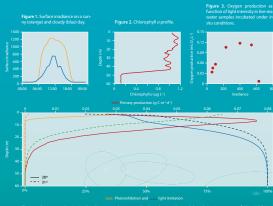


Figure 4. Primary production (red) in the water colum (orange) and light limitation (blue), over a 24 hour pe

Changing ocean conditions

When ocean conditions are changed, by natural or anthropogenic causes it has an effect on the biological pump and the organisms that are part of it.

Increasing atmospheric concentrations of carbon dioxide is reducing the pH of the Increasing atmospheric concentrations of carbon dioxide is reducing the pH of the oceans (by increasing hydrogen ion concentrations), making the sea more acidic. The concentration of calcium carbonate is also decreasing, and this could potentially be detrimental to calcifying organisms, such as corals and some forms of plankton, making them particularly vulnerable to climate change. Because these organisms are part of the biological pump and because CaCO₂ acts as ballast to sinking material, this could potentially have a large inpart on primary production and deep. CoC potentially have a large impact on primary production and deep-sea export of DOM. Warmer temperatures will limit the supply of nutrients in the surface layers, and that will decrease primary production in the euphotic zone. Often, production in surface

layers are limited by nutrient supply from below and an increase in surface layer stratification will have a negative effect on nutrient supply.

Our growing understanding of how our world and seas is working, needs to be passed on to the people who live here, without whom we have no chance of stopping this potentially detrimental effect.

Acknowledgement

This work is part of the marine field course Oceanography at Sea launched by Dansk Center for Havforskning. Associate Professor Colin A. Stedmon (DTU-Aqua), Professor André Viser (DTU-Aqua), Professor Katherine Richardson (Copenhagen University) and Professor Stiig Markager (Aarhus University). Thanks to the Crew at Dana for taking such good care of us!

References



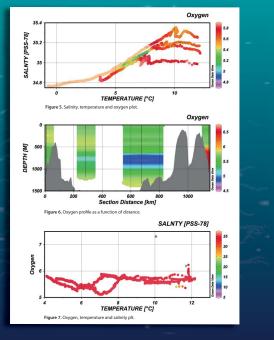
The light regime may change considerably on a small scale time. The surface irradiance measured on a cloudy and sunny day here are only three days apart in

the North Atlantic (fig. 1). Recent windy conditions had likely mixed the upper layer, removing any spa-tial variation with depth but also between stations. There was still stratification however, ensuring a lower layer with few to no algae (fig. 2). As irradiance increases, so does primary production. If the light intensity becomes too high however, the algae will take damage, severely reducing their photosynthesis; this is called photoinhibition (fig. 3).

Based on light profiles from a sunny and cloudy day, incubation experiments at different light intensities and a chlorophyll a profile (to determine the auto-trophic biomass), the primary production at each depth could be calculated over a 24 hour period (fig. 4)

a 24 nour period (ng. 4). Photoinhibition only plays a role in the uppermost layer on a sunny day. Light limitation is most pronounced at depths, but also plays a role in the upper layers due to low light at night. Highest primary production is found on sunny days (1.01 g C m⁻²) compared to cloudy days (0.69 g C m⁻²).

On sunny days, highest primary production is seen at some depth due to photoin-hibition. On cloudy days, irradiance is not sufficient to cause photoinhibition why light is almost always limiting. Here, highest primary production is seen at the surface (fig. 5, 6 and 7).



Estimation of Anthropogenic CO₂ in the

North Atlantic Dansk Center for Havforskning, Dana, summer 2012, Christina Søegren, Nikolaj Thyessen Dam, Sanne Høj Andrén, Stavroula Tsoukali, Thomas Bec

Introduction

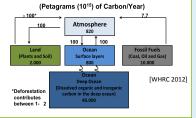
Results

During the course 'Oceanography at sea' on board Dana, we examined different chemical properties at four different station in the North Atlantic. We measured the pH, alkalinity, phosphate and silicate. We used these measurements to estimate the anthropogenic CO₂ in the water column.

ration (µmol/kg)

Global flows of carbon

The ocean plays a dominant role in the Earth's carbon cycle. The figure shows the global carbon cycle, which involves the carbon in and exchanging between the Earth's atmosphere, fossil fuels, the oceans (the surface layers and the deep ocean), the vegetation and soils of the terrestrial ecosystems. The oceans constitute the largest active pool of carbon at the Earth's surface. The total amount of carbon in the ocean is about 50 times greater than the amount in the atmosphere, and is exchanged with the atmosphere on a time-scale of several hundred years.



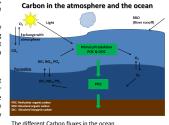
Anthropogenic Carbon calculations

In order to reconstruct the anthropogenic part (C_{ant}) of the observed Dissolved Inorganic Carbon (DIC_{bbc}) in the water samples, it is necessary to derive how much of the carbon/ CO_2 has been removed by primary production (DIC_{bbc}), and how much of the DIC has a natural/preindustrial source (DIC_{pl}).

 $\mathsf{DIC}_{\mathsf{obs}}$ is calculated from the chemical parameters Alkalinity, pH and different physical factors and with an atmospheric $\rm CO_2$ -concentration of 380 ppm. In order to calculate the $\rm DIC_{pl}$ we used the atmospheric CO_2-concentration of 280 ppm, which corresponds to preindustrial values, assuming that the physical characteristics (T, S, nutrients) were the same.

 DIC_{bio} is an estimation of how much CO_2 has been removed by primary production. This is calculated indirectly by estimating how much oxygen has been used by organisms since the last time the water mass was at the surface, under the assumption that the oxygen saturation of surface water is 100%

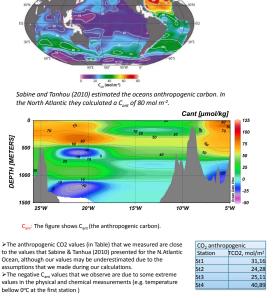
In small timescales DIC_{bio} is insignificant, because the amount of CO_2 that is removed by water is much larger. In a larger timescale however the DIC_{bin} has an important role, because the water bound CO₂ returns to the surface again at some time, but some of the DIC_{bin} will sediment, bury and thereby be removed from the system. [Sarmiento & Gruber, 2006]



 $C_{ant} = DIC_{obs} - \Delta DIC_{bio} - DIC_{pi}$

Depth CO2 SAT% DEPTH [METERS] 100 1500 DIC obs [µ

 25W 20W 25W 15W 10W 10W 10W 10W 10W 10 10 10 Cobstrued), which is higher than the DIC_{pr} The difference between the two is around 55-65 μm kg 1 -



Station	TCO2, mol/m ²
St1	31,16
St2	24,28
St3	25,11
St4	40,89

Conclusion

✓ The silicate and phosphate vertical profiles show a depletion of nutrients at the surface waters, as expected, due to biological activity. \checkmark From our CO₂ %-saturation plot we see that the eastern part of the study area is not saturated yet, which means the

A construction of Co₂ as addition by breastern part of the study are of the

✓ By comparing our results with results in the literature [Sabine & Tanhua 2010, Gruber 1996] we can see that our values are close to already published data.

References

Gruber N., Sarmiento J.L & Stocker T.F. (1996), An improved method for dectectin antropagenic CO2 in the oceans, Global Biogeochemical cycles, vol 10, no. 4 Sabine, C. L. & Tanhua T. (2010), Estimation of Anthropogenic CO2 Inventories in

the Ocean, Annual Reviews. Sarmiento, J. L. & Gruber N. (2006), Ocean Biochemical Dynamics, Princi University Press

Woods Hole Center Resarch Center (WHRC) <u>http://www.whrc.org/global/carbon/index.html</u> (last viewed 14thSeptember 2012)

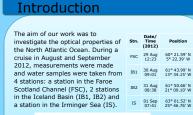


Section of the four stations



Oceanography at Sea: Optics

Ciarán Murray¹, Jakob Thyrring¹, Mette Vodder¹, Ida Ringgaard²



Materials and Methods

I: During CTD casts, light intensity was recorded every 0.20 meters to give profiles of PAR (photosynthetically available radiation) through the water column. The diffuse light attenuation coefficient, K₄, was estimated for the surface waters (0-30m) by linear regression on the log-transformed light intensity

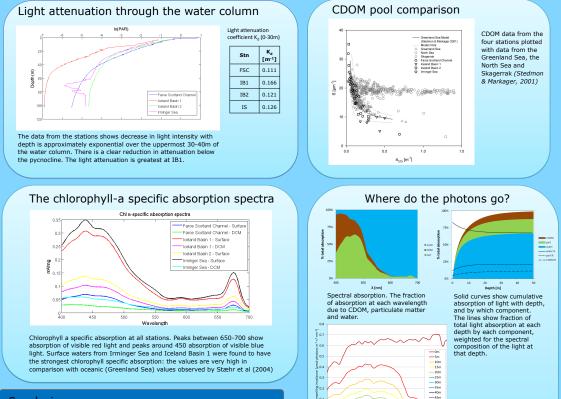
light intensity. III: To estimate chlorophyll specific absorption, water samples from 2 depths (surface and DCM) were filtered onto a Whatman GF/F filter. The optical density of the filters was measured with a Shimadzu UV-2500 spectrophotometer using an integrating sphere attachment. The absorption coefficients were calculated by the following equation: $a(\lambda) = 2.3^{\circ}OD(\lambda)^{*s}/(V_r\beta(\lambda))$ where β is a corrrection factor $\beta(\lambda) = 1.63 * OD(\lambda)^{-0.22}$ (Bricaud & Stramski 1990)

To calculate the chlorophyll specific absorption, the absorption coefficients were divided by the measured chlorophyll-a concentrations at the respective depths.

childrophylin-a concentrations at the respective depths. III: to investigate the origin of CDOM, water samples were collected from each station. Water was filtered through a 0.2 µm filter, and by using 10 cm quartz cuvettes the samples were analysed in a spectrophotometer. An exponential function was fitted to the observed absorption spectra to give estimates of spectral slope coefficient (5) and absorption coefficient at 375 nm (a₃₇₅): $a(\lambda)=a(\lambda_0)*exp(S(\lambda_0-\lambda))$ (Stedmon & Markager, 2001)

W: A photon budget was made for station FSC. Spectral attenuation was modelled by the sum of calculated absorption components and backscattering $K_d(\lambda)=(a_{mater}(\lambda)+a_{CDOM}(\lambda)+a_{part}(\lambda)+b_p)/cos(\theta)$. The backscatter b_b and zenith angle θ were estimated by fitting the model to the observed PAR profile.

Results and Discussion



Conclusions

PAR attenuatedthrough the water column and the highest attenuation was seen at Iceland Basin 1. There was no clear relation between chlorophyll-a specific absorption and depth or

The CDOM seen in the four stations had similar optical charateristics to CDOM from the Greenland sea and most likely derives from in situ production in the ocean.

References Marcuick, A. and D. Stramski. 1990. Spectral Absorption Coefficients of Living Phytoplankton and Nonalgal Biogenous Matter. Limnol. Oceaogr. 35: 562-582 Stadmon, CA. and S. Markagor. 2001. The Optics of Chromophoric Dissolved Organic Matter (CDOM) in the Greenlat Sea. Immol. Oceaogr. **46**: 2087-2093 Stabir et al. 2004. Pigment speacific in vivo light absorption of phytoplankton from estuarine, coastal and oceanic watt Mar. Ecol. Prog. Ser. **275**: 115-128

Modelled irradiance spectra by depth.

