











SECOND INTERNATIONAL SYMPOSIUM ON THE OCEAN IN A HIGH-CO₂ WORLD

MONACO - OCTOBER 6-9, 2008

BOOK OF ABSTRACTS AND PROGRAMME













Front Cover Photos - credits from left to right:

Sea-bream fish eggs (© Jean-Louis Teyssié)

Floating open-ocean mesocosms (© Ulf Riebesell)

Cold-water coral *Lophelia pertusa* (©André Freiwald)

Cold-water coral reef (© Karen Hissmann)

Sea-bream fish larvae (© Jean-Louis Teyssié)

Increasing attention is being given to the carbon footprint of travel, including that of scientists attending meetings. The international Planning Committee for this symposium has agreed to make a US\$5,000 contribution to a reputable carbon offset organization, Climate Care (http://www.climatecare.org/). This contribution will at least partially offset the carbon emitted by the travel of participants in this symposium. The sponsors understand that carbon offsets are an imperfect substitute for reduced travel.

SECOND INTERNATIONAL SYMPOSIUM ON

THE OCEAN IN A HIGH-CO₂ WORLD

The scientific sponsors and the organizing committees of the Symposium gratefully acknowledge the financial and in-kind support received from the following organizations and funding agencies:

U.S. National Science Foundation
Prince Albert II of Monaco Foundation
Intergovernmental Oceanographic Commission of UNESCO
The Scientific Committee on Oceanic Research
International Atomic Energy Agency
International Geosphere-Biosphere Programme
Musée Oceanographique
Centre Scientifique de Monaco

Second Symposium on the Ocean in a High-CO₂ World Book of Abstracts and Programme FOREWORD

When we met at UNESCO headquarters in Paris four years ago at the 1st Symposium on the Ocean in a High-CO2 World, about 120 scientists gathered to discuss how recent increases in atmospheric CO2 were affecting ocean chemistry and biology, to evaluate potential strategies to artificially enhance ocean carbon uptake, and to discuss directions for future research. The term ocean acidification was not in wide use, and only a small group of specialists had been studying how increasing marine concentrations of CO2 and corresponding reductions in pH and carbonate ion concentrations were affecting marine organisms, mostly corals. For years, ocean scientists had thought mostly about the beneficial effects of the ocean's great capacity to take up CO2, thereby moderating the increase in atmospheric CO2 from fossil-fuel combustion. But as the meeting progressed, there was a growing awareness of many problems associated with corresponding changes in ocean chemistry and associated biological impacts. That 1st symposium marked a turning point for many of us. We suddenly understood that impacts were as important as air-sea CO2 fluxes. The media also picked up on these heightened scientific concerns and further increased their fervor when subsequent national reports were released and when three papers presented at the symposium were published in high-profile science journals.

Four years later, we meet again under heightened concern. There is now much wider scientific interest, as illustrated by the 250 scientists from 32 countries who are attending this 2nd symposium, many of whom are now directly involved in ocean-acidification research. Ocean acidification is now widely cited in the press in conjunction with climate change, often being referred to as *the other CO*₂ *problem*. The convenors of the 1st symposium, IOC-UNESCO and SCOR, have been joined by two other international bodies, the IAEA and IGBP. There has been progress to begin to fund efforts to understand the fundamental science concerns, impacts on key organisms, and how those might affect ecosystems in general. There is now a political urgency to communicate new science findings to economists and policymakers, to educate the public, and to train future scientists. Our 2nd symposium provides a small step in this direction by following up the 3-day science meeting with a 1-day wrapup session where scientists will meet with both economists and policymakers, in the morning, and secondary-school science teachers, in the afternoon.

It is no accident that the 2nd Symposium on the Ocean in a High-CO₂ World is being held in Monaco at the Musée Océanographique, which was founded nearly 100 years ago by Prince Albert I, who was himself an oceanographer and whose marine collections are still on display. It is hard to imagine a more conducive environment for you to discuss advances since the 1st symposium, to plan future high-priority research, and to begin to build the critical links among scientists, economists, and policymakers that our society requires in order to realize the full extent of the global and long-term impacts of ocean acidification and to motivate it to take action.

James C. Orr, Chairman, International Scientific Planning Committee

Second Symposium on the Ocean in a High-CO₂ World Book of Abstracts and Programme

TABLE OF CONTENTS

FORWARD	i
PLANNING COMMITTEES	1
LOCAL INFORMATION	3
MAPS	3
PUBLIC TRANSPORTATION IN MONACO	4
BUS ROUTE MAP	5
SUMMARY PROGRAMME	7
DETAILED DAILY PROGRAMME	
DAY 1 - MONDAY, OCTOBER 6	9
DAY 2 - TUESDAY, OCTOBER 7	10
DAY 3 - WEDNESDAY, OCTOBER 8	12
DAY 4 - THURSDAY, OCTOBER 9	13
SPEAKERS' ABSTRACTS	
DAY 1 - MONDAY, OCTOBER 6	15
DAY 2 - TUESDAY, OCTOBER 7	24
DAY 3 - WEDNESDAY, OCTOBER 8	31
DAY 4 - THURSDAY, OCTOBER 9	35
POSTER ABSTRACTS	37
LIST OF PARTICIPANTS	91
AUTHOR INDEX	107

Second Symposium on the Ocean in a High-CO2 World

PLANNING COMMITTEES

International Scientific Planning Committee:

Ken Caldeira Victoria Fabry André Freiwald Jean-Pierre Gattuso Peter Haugan Patrick Lehodey James Orr (Chair) Silvio Pantoja Hans-O. Pörtner Ulf Riebesell Tom Trull

Local Scientific Organizing Committee:

Denis Allemand (Centre Scientifique de Monaco)
Michel Boisson (Centre Scientifique de Monaco)
Jean-Pierre Gattuso (Laboratoire Océanographique de Villefranche)
Nadia Ounais (Musée Océanographique, Monaco)
Stéphanie Reynaud (Centre Scientifique de Monaco)
James Orr (Marine Environment Laboratoires/IAEA, Monaco)
Philippe Mondielli (Fondation Albert II)

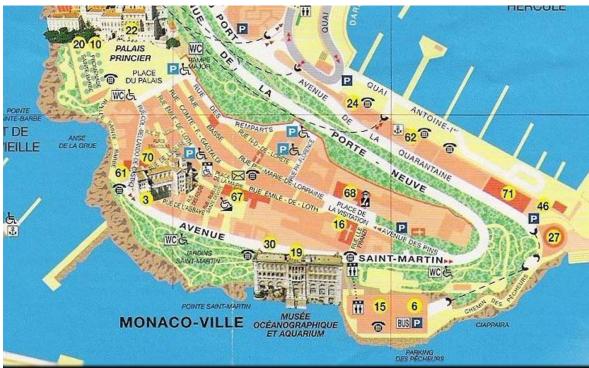
Staff Members from Sponsoring Organizations:

Elizabeth Gross, SCOR, Symposium Manager Wendy Broadgate, IGBP Maria Hood, IOC Edward R. Urban, Jr., SCOR Mary Ann Williams, IGBP Sue Williams, UNESCO

Monaco - Local Information

The schematic maps below will give you a general idea of the areas of Monaco. The lower one gives more detail about "Monaco-ville", the old town close to the Musée Océanographique. We hope to have more detailed maps and a restaurant list available at the registration desk, and your hotel may also provide them.





Public Transportation in Monaco:

As in most European countries, using public transportation is efficient. Although walking distances are short - the whole country is less than 500 acres - you may wish to use the bus to get from your hotel to the Musée Océanographique. The following information is taken from the Web site of the local bus company:

Description of the Bus Routes:

```
5 bus lines to get you where you are going...
```

```
Line 1: Monaco Ville - Casino - Saint Roman (Round Trip).
```

Line 2: Monaco Ville - Casino - Jardin Exotique (Round Trip).

Line 4: Gare - Larvotto (Plages) (Round Trip).

Line 5 : Gare -Fontvieille - Hôpital - Gare.

Line 6: Fontvieille - Casino - Larvotto (Plages) (Round Trip).

How to get a Ticket:

In the bus...

Next to the conductor, upon boarding the bus. (Single trip tickets, 8 trip card, 8 trip "Monaco - Beausoleil", 4 trip card and daily tourist card).

At our offices...

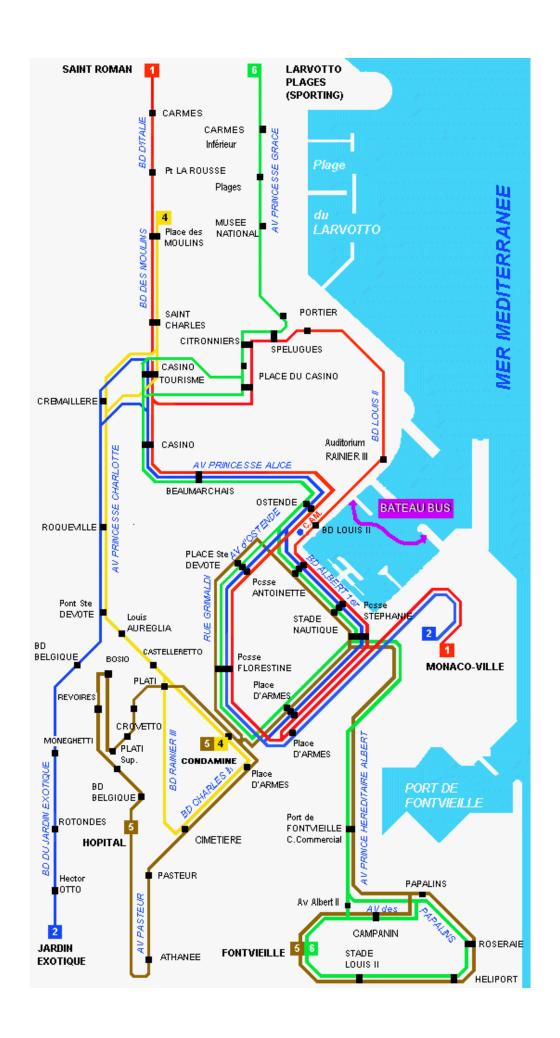
Compagnie des Autobus de Monaco, 3, av. du président J.F. Kennedy, 98000 Monaco. (Only for tickets not sold on the buses)

Riding the Bus:

Go to one of the 141 bus stops of the Principality's network... Upon boarding the bus...

- 1. Boarding the bus is done exclusively by the front unless authorized otherwise by the conductor or ticket inspector of the C.A.M..
- 2. Get your ticket or ask the conductor.
- 3. Validate your ticket in the machine provided for that purpose, then keep the ticket in your possession during the rest of the trip.

A map of the bus routes is shown on the next page.



Second Symposium on the Ocean in a High-CO₂ World Summary Programme

SUNDAY, 5 OCTOER	MONDAY, 6 OCTOBER	TUESDAY, 7 OCTOBER	WEDNESDAY, 8 OCTOBER	THURSDAY, 9 OCTOBER
	Registration 8:00-9:00	First session begins 8:30	First session begins 8:30	Beyond natural science Opening by HSH Prince Albert II 9:00
	Opening 9:00-9:30	Mechanisms of Calcification	Adaptation and microevolution	ns Economics ical and Policy
			New concerns	
	Scenarios of ocean acidification	Physiological effects: from microbes to fish	Biogeochemical consequences and feedbacks	
	Coffee Break 10:45-11:15	Coffee Break 10:30-11:00	Coffee Break 10:15-10:45	Coffee Break 10:50-11:20
	Impacts on benthic and pelagic calcifiers	Physiological effects: from microbes to fish continued	Biogeochemical consequences and feedbacks continued	Press Conference 11:30-12:30
	Lunch 12:00 - 14:00	Lunch 12:00 - 14:00	Lunch 11:30-13:00	
	Impacts on benthic and pelagic calcifiers continued	Fisheries, food webs, and ecosystem impacts	Breakout Sessions	
	Coffee Break 15:15 - 15:45	Coffee Break 15:30 - 16:00	Coffee Break 15:15 - 15:45	Outreach for local students, teachers 14:00-16:00
	Ocean carbon system: past & present	CO ₂ Disposal	Reports from Breakout Sessions	
	Effects of ocean acidification on nutrient and metal speciation	Poster Session 2 17:00-19:00	Closing summaries 17:00-18:00	
Registration Opens 17:00	Poster Session 1 With refreshments 18:00-20:00	Cocktail reception 19:00-20:00		
Ice-Breaker and registration 18:00-19:30		Dinner 20:00-22:00		

Second Symposium on the Ocean in a High-CO₂ World Detailed Programme

SUNDAY, 5 OCTOBER

Ice-Breaker Reception (and Registration)

	Tee Diemer Heepman (untu Hegasin unteri)			
17:00 – 18:00	Registration (access to the Musée Océanographique after registering)			
18:00 – 19:30	Ice-Breaker Reception (registration continues; access to 1st floor access only)			
DAY 1: MONDAY, 6 OCTOBER				
8:00 – 9:00	Registration			
9:00 - 9:30 9:00 - 9:15	Opening & Welcome Robert Calcagno – Minister for Public Works, the Environment and Urban Development (Monaco)			
9:15 - 9:20 9:20 - 9:30	Representative of the Musée Océanographique James Orr – MEL/IAEA, Monaco, Chair, Planning Committees			
Scenarios of ocean acidification – James Orr, Chair				
9:30 – 10:00	<i>Invited:</i> Present and future changes of carbonate systems in the global oceans – Richard Feely , NOAA/PMEL, Seattle, USA			
10:00 - 10:15	Impact of climate change mitigation on ocean acidification projections – Gian-Kasper Plattner , ETH Zurich, Switzerland			
10:15 – 10:30	CO ₂ emission targets for future changes in ocean carbon chemistry – Richard Zeebe , University of Hawaii, USA			
10:30 – 10:45	High vulnerability of Eastern boundary upwelling systems to ocean acidification – Nicolas Gruber , ETH Zurich, Switzerland			
10:45 – 11:15	Coffee Break			
Impacts on benthic and pelagic calcifiers – Denis Allemand, Chair				
11:15 – 11:45	<i>Invited:</i> Impact of ocean acidification on benthic organisms – Jean-Pierre Gattuso , LOV, Villefranche-sur-mer, France			
11:45 – 12:00	Poorly cemented coral reefs of the eastern tropical Pacific: possible insights into reef development in a high-CO ₂ world– Joan Kleypas , National Center for Atmospheric Research, USA			
12:00 – 14:00	Lunch			
14:00 – 14:15	The impact of ocean acidification and temperature on the reproduction and			

development of oysters and the potential of genetic differences to ameliorate climate change – Laura Parker, University of Western Sydney, Australia

14:15 – 14:30 Latitudinal variation in calcification; vulnerability of Antarctic benthic calcifiers to ocean acidification – **Sue-Ann Watson**, National Oceanography Centre Southampton, UK 14:30 – 15:00 *Invited:* Pelagic calcifiers: pteropods and forams – **Victoria Fabry**, Cal. State University, San Marcos, USA Interannual variability of pteropod shell weights in the high-CO₂ Southern Ocean – 15:00 - 15:15Donna Roberts, Antarctic Climate & Ecosystems CRC, Hobart, Australia 15.15 Coffee break Ocean carbon system: past & present – Peter Haugan, Chair 15:45 – 16:15 *Invited*: Controls on evolution of ocean carbonate chemistry over the past 10⁹ years – Ken Caldeira, Department of Global Ecology, Carnegie Institution, Stanford, USA 16:15 – 16:30 Boron isotope evidence of ocean acidification in the Neoproterozoic – **Simone** Kasemann, University of Edinburgh, UK 16:30 - 16:45Reduced calcification in modern Southern Ocean planktonic foraminifera – Andrew Moy (William Howard), Antarctic Climate & Ecosystems CRC, University of Tasmania, Australia 16:45 – 17:00 Current rates of change in pH and calcium carbonate saturation in the high latitude North Atlantic Ocean - Jon Olafsson, Marine Research Institute, Reykjavik, Iceland Low winter CaCO₃ saturation in the Baltic Sea and consequences for calcifiers – 17:00 - 17:15**Toby Tyrrell**, National Oceanography Centre, Southampton University, UK Effects of ocean acidification on nutrient and metal speciation -Silvio Pantoja, Chair 17:15 – 17:45 *Invited:* Ocean acidification and metal speciation – **Hein de Baar**, Royal Netherlands Institute for Sea Research, the Netherlands 17:45 – 18:00 Ocean acidification effects on iron speciation in seawater – **Eike Breitbarth**, Department of Chemistry, University of Gothenburg, Sweden 18:00 – 20:00 **Poster Session 1** (with refreshments) DAY 2: TUESDAY, 7 OCTOBER Mechanisms of Calcification - Joan Kleypas, Chair 8:30 - 9:00Invited: Biomineralization mechanisms in marine calcifiers in view of ocean acidification - Jonathan Erez, Institute of Earth Sciences, Hebrew University of Jerusalem, Israel 9:00 - 9:15Effect of acidification on coral calcification: working hypothesis towards a physiological mechanism – Francesca Marubini (Denis Allemand), Centre Scientifique de Monaco

9:15 – 9:30 Predictions of carbon fixation during a bloom of *Emiliyana Huxleyi* is highly sensitive to assumed response to shift in pCO₂ – **Olivier Bernard**, INRIA-COMORE, Sophia-Antipolis, France

Physiological effects: from microbes to fish - Victoria Fabry, Chair

- 9:30 10:00 *Invited:* Physiological Mechanisms Linking Climate to Ecosystem Change: Effects of Ocean Acidification on Marine Animals in Times of Ocean Warming **Hans-Otto Pörtner**, Alfred Wegener Institute for Polar and Marine Research (AWI), Germany
- 10:00 10:30 *Invited:* Impacts of Ocean Change on Primary Producers **Ulf Riebesell,** Leibniz Institute of Marine Sciences (IFM-GEOMAR), Germany
- 10:30 11:00 **Coffee Break**
- 11:00 11:30 *Invited:* Physiology overview of Microbes **Antje Boetius**, MPI für Marine Microbiology, Bremen, Germany
- 11:30 11:45 Effects of hypercapnic acidification of seawater on the biology of non-calcifying marine organisms **Erik Thuesen**, Evergreen State College, Laboratory, Olympia, USA
- 11:45 12:00 Predicting the impact of ocean acidification on benthic biodiversity: what can animal physiology tell us? **Stephen Widdicombe**, Plymouth Marine Laboratory, UK
- 12:00 14:00 *Lunch*

Fisheries, food webs, and ecosystem impacts (PICES-ICES session) – Patrick Lehodey, Chair

- 14:00 14:30 *Invited:* Consequences of Ocean Acidification for Fisheries **Jan Helge Fosså**, Institute of Marine Research, Bergen, Norway
- 14:30 15:00 An ocean acidification simulation experiment with benthic animals using a precise pCO2 control system **Yukihiro Nojiri**, CGER/NIES, Tskuba, Japan
- 15:00 15:15 Natural CO2 vents reveal ecological tipping points due to ocean acidification **Jason Hall-Spencer**, Marine Institute, Biological Sciences, University of Plymouth, UK
- 15:15 15:30 Salmon pHishing in the northeast Pacific; an archaeological dig in the North Pacific survey data (1956–1964) **Skip McKinnell**, North Pacific Marine Science Organization, Sidney, Canada
- 15:30 16:00 *Coffee Break*

CO₂ Disposal – Ken Caldeira, Chair

- 16:00 16:30 *Invited:* Effects of CO₂ capture and storage on ocean acidification **Peter Haugan**, Geophysical Institute, University of Bergen, Norway
- 16:30 16:45 Modelling of CO₂ dispersion leaked from seafloor off Japanese coast **Yuki Kano** (**Toru Sato**), AIST (and University of Tokyo), Japan

Tuesday Evening Symposium Events:

- 19:00 20:00 Symposium Cocktail Aquarium (in the Musée Océanographique)
- 20:00 22:00 Symposium Dinner 1st Floor of Musée Océanographique (stand-up buffet)

DAY 3: WEDNESDAY, 8 OCTOBER

Adaptation and microevolution - Ulf Riebesell, Chair

- 8:30 9:00 *Invited:* A brief history of skeletons in the ocean **Andrew Knoll**, Harvard University, USA
- 9:00 9:15 Influence of high CO₂ on coccolithophores under long-term cultivation **Marius Müller**, Leibniz Institute of Marine Sciences, Germany

New concerns - Ulf Riebesell, Chair

- 9:15 9:30 Impact of ocean acidification on underwater sound: reduced low frequency absorption, increased noise levels, potentially higher stress for marine mammals **David Browning**, Department of Physics, University of Rhode Island, USA
- 9:30 9:45 Experimental approaches of carbonate chemistry manipulation in CO₂ perturbation studies **Kai Schulz**, Leibniz Institute for Marine Sciences (IFM-GEOMAR), Germany

Biogeochemical consequences and feedbacks to the Earth system – Tom Trull, Chair

- 9:45 10:15 *Invited :* Biogeochemical consequences of ocean acidification **Laurent Bopp**, Laboratoire des Sciences du Climat et de l'Environnement CEA-CNRS-UVSQ, IPSL, Gif-sur-Yvette, France
- 10:15 10:45 *Coffee Break*
- 10:45 11:00 Dissolution of CaCO₃ in shallow water carbonate environments in the high CO₂ world of the Anthropocene **Andreas Andersson**, Bermuda Institute of Ocean Sciences, Bermuda
- 11:00 11:15 Impacts of ocean acidification on marine biogenic trace gas production **Frances Hopkins**, Laboratory for Global Marine and Atmospheric Chemistry, School of
 Environmental Sciences, University of East Anglia, Norwich, UK
- 11:15–11:30 From laboratory manipulations to Earth system models: an 'Eppley curve' for calcification rate? **Andy Ridgwell**, University of Bristol, UK
- 11:30 13:00 *Lunch*

13:00 – 15:15 **Breakout sessions**

- Natural and Artificial Perturbation Experiments to Assess Acidification (e.g., paleoceanography, spatial variability, and mesocosm studies, modeling)
 Chair: Ulf Riebesell Rapporteur: Steve Widdicombe
- Observational Networks for Tracking Acidification and its Impacts (e.g., sensor development, observation networks, ecosystem responses, modelling)
 Chair: Toby Tyrrell Rapporteur: Chris Sabine
- 3. Scaling Organism to Ecosystem Acidification Effects and Feedbacks on Climate (e.g., organism dose-response, modeling)

Chair: Hans-Otto Pörtner Rapporteur: Ken Caldeira

- 15:15 15:45 *Coffee Break*
- 15:45 17:00 Reports from Breakout sessions

Closing summaries for 3-day science meeting – James Orr, Chair

- 17:00 17:30 What have we learnt about ocean acidification since 2004 and where should we be in 2012? **Jean-Pierre Gattuso -** LOV, Villefranche-sur-mer, France Followed by discussion.
- 17:30 18:00 Closing statement James Orr, Planning committee

DAY 4: THURSDAY, 9 OCTOBER

Beyond natural science

- 09:00 09:30 Science summary from first three days of the symposium **Carol Turley**, Plymouth Marine Laboratory, UK
- 09:30 10:00 Basic Economics of Ocean Acidification **Hermann Held**, Potsdam Institute for Climate Impact Resrearch, Germany
- 10:00 10:30 Ocean acidification connecting the science to policy **John Baxter**, Scottish Natural Heritage, UK
- 10:30 10:50 Opening H.S.H. Prince Albert II, Monaco
- 10:50 11:20 *Coffee Break*

Press conference

- 11:30 12:30 Press conference in English and French
- 12:30 13:30 *Lunch*

Outreach for local students and teachers (in French)

14:00 – 16:00 Presentations and hands-on workshop potentially organized with CarboSchools, EPOCA EU Project, Musée Océanographique, Océanopolis, & l'Education Nationale de Monaco

SPEAKERS' ABSTRACTS

The abstracts in this section are arranged in order of their appearance in the Symposium programme.

I = Invited Speaker C = Contributed Talk

DAY 1 - MONDAY, OCTOBER 6 2008

SCENARIOS OF OCEAN ACIDIFICATION

PRESENT AND FUTURE CHANGES OF THE CARBONATE SYSTEM IN THE GLOBAL OCEANS [I]

<u>Feely, Richard A</u>¹, Christopher L. Sabine¹, James C. Orr², Robert H. Byrne³, Frank J. Millero⁴ and Dana Greeley¹

The addition of fossil fuel carbon dioxide to the atmosphere is rapidly changing seawater chemistry and the calcium carbonate saturation state of the world's oceans as a result of the acidifying effects of CO₂ on seawater. This acidification makes it more difficult for marine organisms (e.g., corals, plankton, calcareous algae, and mollusks) to build skeletons, tests, and shells of calcium carbonate. Impacts on these calcifying organisms will lead to cascading effects throughout marine ecosystems. Repeat hydrographic cruises and modeling studies in the Atlantic, Pacific and Indian Oceans show evidence for increased ocean acidification. The dissolved inorganic carbon increases, of about 10-15 µmol kg⁻¹ in surface and intermediate waters over the past 15 years, are consistent with corresponding pH decreases of approximately 0.025 units in surface waters. These dramatic changes can be attributed, in most part, to anthropogenic CO₂ uptake by the ocean. These data verify earlier model projections that the oceans are undergoing ocean acidification as a result of the uptake of carbon dioxide released as a result of the burning of fossil fuels. From these results we have estimated an average upward migration of the aragonite saturation horizon of approximately 1-2 m yr⁻¹ in the Pacific and Indian Oceans. Such shoaling is due to the effects of anthropogenic CO₂, ventilation and biological respiration processes in the surface and intermediate waters.

IMPACT OF CLIMATE CHANGE MITIGATION ON OCEAN ACIDIFICATION PROJECTIONS [C]

<u>Plattner, Gian-Kasper</u>¹, Fortunat Joos^{2,3}, Kuno M. Strassmann², Marco Steinacher², and Nicolas Gruber¹

Ocean acidification due to rising atmospheric CO₂ concentrations lowers ocean pH and CaCO₃ saturation and might eventually lead to undersaturation and dissolution of CaCO₃ shells in parts of the ocean with potentially negative consequences for marine ecosystems. The magnitude and timing of changes in surface pH and CaCO₃ saturation state are largely driven by atmospheric CO₂ levels and thus critically depend on the amount of future CO₂ emissions. Here we investigate the effect of

¹Pacific Marine Environmental Laboratory, National Oceanic and Atmospheric Administration, Seattle, Washington 98115, USA (Richard.A.Feely@noaa.gov).

²Marine Environment Laboratory, IAEA, 4 Quai Antoine 1^{er}, MC-98000, Monaco.

³College of Marine Science, University of South Florida, 140 Seventh Avenue South, St. Petersburg, FL 33701, USA

⁴University of Miami/Rosensteil School of Marine and Atmospheric Sciences, Miami, FL 33149, USA

climate mitigation actions on ocean acidification projected with the Bern2.5CC carbon cycle-climate model for a set of newly-developed multi-gas reference and climate mitigation emissions scenarios from Integrated Assessment Models. These scenarios cover a wide range of plausible 21st century emissions based on projected evolutions of the socio-economic-technological system. Reduced CO₂ emissions in climate mitigation scenarios compared to reference scenarios lead to lower projected atmospheric CO₂ concentrations and to smaller reductions in surface pH and CO₃²-ion concentrations. In addition, climate mitigation also results in a substantial delay in the timing when critical thresholds will be passed, e.g. when the surface ocean becomes undersaturated with regard to CaCO₃. We will quantify these climate mitigation-induced differences in ocean acidification projections, both in terms of absolute values as well as in terms of temporal evolution. The effects of climate sensitivity and carbon cycle related uncertainties on projected ocean acidification will also be discussed, highlighting the substantial uncertainties associated with these projections.

${\rm CO_2}$ EMISSION TAGETS FOR TOLERABLE FUTURE CHANGES IN OCEAN CARBONATE CHEMISTRY [C]

Zeebe, Richard E.¹, James C. Zachos², Ken Caldeira³, and Toby Tyrrell⁴

The extent of acidification and pH decline in the surface ocean is controlled by the total amount and release time of anthropogenic carbon dioxide. In turn, these variables will determine the degree of detrimental consequences for marine organisms. This includes possible reduction in calcification rates in coccolithophores, foraminifera, and corals, as well as potential negative implications for coral reef communities and shore protection. The allowable drop in surface ocean pH, which would prevent such effects, is not yet known because of our incomplete understanding of the physiological response of the various marine organisms. Intensified research efforts are therefore required to close this gap. Nevertheless, for a prescribed target decline in future surface ocean pH, the corresponding total amount and release time of anthropogenic carbon can be calculated. Fortunately, and in contrast to climate model predictions, such future ocean chemistry projections are largely model-independent on a timescale of a few centuries, mainly because the chemistry of CO₂ in seawater is well known. We present results from a coupled carbon cycle model (including deep-sea sediments) to address this issue. Our results provide specific constraints on future CO₂ emissions, if a shift in surface ocean pH and carbonate saturation state beyond a given threshold is to be avoided. Our projected changes in ocean carbonate chemistry should serve as a guideline for policy protocols that identify CO₂ emission targets to reduce the effects of man-made ocean acidification.

HIGH VULNERABILITY OF EASTERN BOUNDARY UPWELLING SYSTEMS TO OCEAN ACIDIFICATION [C]

Gruber, Nicolas¹, Hartmut Frenzel², Claudine Hauri¹, Zouhair Lachkar¹, and Gian-Kasper Plattner¹

Eastern Boundary Upwelling Systems (EBUS), such as the California or Canary Current Systems are particularly sensitive to ocean acidification as their pH is already low and their change in pH for a given uptake of anthropogenic CO₂ is particularly high. This is mostly a consequence of the upwelling of waters with low pH and high concentrations of dissolved inorganic carbon in the absence of high

¹ Environmental Physics, Institute of Biogeochemistry and Pollutant Dynamics, ETH Zurich, Zurich, Switzerland (gian-kasper.plattner@env.ethz.ch)

²Climate and Environmental Physics, Physics Institute, University of Bern, Bern, Switzerland

³ Oeschger Centre for Climate Change Research, University of Bern, Bern, Switzerland

¹School of Ocean and Earth Science and Technology, Dept of Oceanography, University of Hawaii, 1000 Pope Road, MSB 504, Honolulu, HI 96822, USA. (zeebe@hawaii.edu)

²Earth and Planetary Sciences Dept., UCSC, Santa Cruz, CA 95060, USA.

³Department of Global Ecology, Carnegie Institution, Stanford, CA 94305, USA. ⁴National Oceanography Centre, Southampton University, European Way, Southampton, SO14 3ZH, U.K.

alkalinity, i.e. waters with very low carbonate ion concentrations. This effect is particularly strong in the Pacific Ocean, as evidenced by the dramatic shoaling of the aragonite saturation horizon off the U.S. West Coast to depths of less than 100 m as recently reported in the literature. We investigate this high vulnerability to ocean acidification of EBUS by conducting eddy-resolving simulations with the Regional Oceanic Modeling System (ROMS) coupled to a state-of-the art ecosystem model for three of the four major EBUS, namely the California, the Canary, and the Humboldt Current Systems. We will investigate the past, present and future changes in pH, CaCO₃ saturation horizons, and other biogeochemical and ecological processes in response to elevated atmospheric CO₂ using the observed atmospheric CO₂ record over the historical period and prescribing a business as usual emission scenario over the 21st century. A particular focus of our analyses is on the rate of change and on the timing when critical thresholds will be passed in the different EBUS, e.g. when the aragonite saturation horizon will enter the euphotic zone.

IMPACTS ON BENTHIC AND PELAGIC CALCIFIERS

IMPACT OF OCEAN ACIDIFICATION ON BENTHIC ORGANISMS [I] Gattuso, Jean-Pierre¹

The response of marine organisms to ocean acidification was first investigated using perturbation experiments on benthic organisms (tropical coralline algae and corals). At the time of the 1st meeting on "The Oceans in a High-CO2 World" (2004), very little information on the response of benthic organisms to environmentally relevant pH or pCO2 levels was available, except on corals and, to a lesser extent, echinoderms. Only one study had examined the interaction between elevated pCO2 and temperature. Calcification and net primary production were, by far, the main processes investigated. This presentation will briefly summarize the current knowledge on the response of benthic organisms, highlight the progress made since 2004 and will point to research gaps. The new studies generally confirm the overwhelming evidence that calcification of almost all organisms assayed decreases at low pH/high pCO2. Among the new information now available is the effects on other processes, mostly physiological processes such as immune responses, new or additional organisms (seagrasses, molluscs, echinoderms, benthic foraminifera), larval stages (molluscs), and ecological processes such as recruitment (coralline algae) and predator-prey relationships (between a mollusk and a macroalga). An innovative development is the recent use of natural changes in the carbonate chemistry, resulting from diurnal patterns or occurring in areas naturally enriched in CO2, to investigate the impact of ocean acidification at organism and community levels. Calcification and dissolution of CaCO3 remain the processes the investigated the most and very few other processes have been considered.

POORLY CEMENTED CORAL REEFS OF THE EASTERN TROPICAL PACIFIC: POSSIBLE INSIGHTS INTO REEF DEVELOPMENT IN A HIGH-CO₂ WORLD [C]

Manzello, Derek P.¹, <u>Joan A. Kleypas</u>², David A. Budd³, C. Mark Eakin⁴, Peter W. Glynn⁵, and Chris Langdon⁵

Ocean acidification describes the progressive, global reduction in seawater pH that is currently underway due to the accelerating oceanic uptake of atmospheric CO₂. Acidification is expected to reduce coral reef calcification and increase reef dissolution. Inorganic cementation in reefs describes the precipitation of CaCO₃ that acts to bind framework components and occlude porosity. Little is known about the effects of ocean acidification on reef cementation and whether changes in

¹ Environmental Physics, Institute of Biogeochemistry and Pollutant Dynamics, ETH Zurich, Zurich, Switzerland (nicolas.gruber@env.ethz.ch)

² IGPP, University of California, Los Angeles, Los Angeles, CA.

¹ Laboratoire d'Océanographie, CNRS-Université Pierre et Marie Curie-Paris 6, B. P. 28, 06234 Villefranche-sur-mer Cedex, France (gattuso@obs-vlfr.fr)

cementation rates will affect reef resistance to erosion. Coral reefs of the eastern tropical Pacific (ETP) are poorly developed and subject to rapid bioerosion. Upwelling processes mix cool, subthermocline waters with elevated pCO₂ (the partial pressure of CO₂) and nutrients into the surface layers throughout the ETP. Concerns about ocean acidification have led to the suggestion that this region of naturally low pH waters may serve as a model of coral reef development in a high-CO₂ world. We analyzed seawater chemistry and reef framework samples from multiple reef sites in the ETP and found that a low carbonate saturation state (Ω) and trace abundances of cement are characteristic of these reefs. These low cement abundances may be a factor in the high bioerosion rates previously reported for ETP reefs, although elevated nutrients in upwelled waters may also be limiting cementation and/or stimulating bioerosion. ETP reefs represent a real-world example of coral reef growth in low- Ω waters that provide insights into how the biological-geological interface of coral reef ecosystems will change in a high-CO₂ world.

THE IMPACT OF OCEAN ACIDIFICATION AND TEMPERATURE ON THE REPRODUCTION AND DEVELOPMENT OF OYSTERS AND THE POTENTIAL OF GENETIC DIFFERENCES TO AMELIORATE CLIMATE CHANGE [C]

Parker, Laura M.¹, Pauline M Ross¹ and Wayne A. O'Connor²

Studies have found that elevations in atmospheric carbon dioxide (CO₂), predicted as early as 2065, are expected to have impacts on organisms in oceanic environments. While studies have investigated the effect of elevated pCO₂ on the calcification of adult marine organisms, less is known about the effect of elevated pCO₂ and temperature on the early life history stages nor their synergistic effect We examined the synergistic effect of elevated pCO_2 (375, 600, 750 and 1000 ppm) and temperature (18, 22, 26 and 30°C) on the fertilisation and embryonic development of the Sydney Rock Oyster Saccostrea glomerata (Gould, 1850) and the Pacific Oyster Crassostrea gigas and determined whether there were genetic differences which may have the potential to ameliorate the impacts of climate change. Overall as pCO_2 increased, fertilisation decreased. The temperature of 26°C was the optimum temperature for fertilisation, as temperature increased and decreased from this optimum, fertilisation decreased. There was also an effect of pCO_2 and temperature on embryonic development. Generally as pCO2 increased, the percentage and size of D-veligers decreased and the percentage of D-veligers that were abnormal increased. S. glomerata spat from mass selected and pair mated fast growth lines were reared at ambient (present concentration: 375 ppm) and elevated CO₂ (1000 ppm). There were significant differences in the growth of selected lines of S. glomerata between elevated and ambient CO₂. This research provides preliminary data that selective breeding of S. glomerata and other estuarine molluses, maybe a potential strategy to overcoming impacts of climate change.

¹ Cooperative Institute of Marine and Atmospheric Studies, Rosenstiel School, Marine Biology and Fisheries, Univ. of Miami, 4600 Rickenbacker Cswy., Miami, FL 33149, USA

² Inst for the Study of Society and Environment, National Center for Atmospheric Research, Boulder CO 80307, USA (kleypas@ucar.edu)

³ Dept of Geological Sciences, Univ. of Colorado, Boulder CO 80309, USA

⁴ National Oceanic and Atmospheric Administration, Coral Reef Watch, 1335, East-West Highway, Silver Spring, MD 20910 USA

⁵ Rosenstiel School, Univ. of Miami, Marine Biology and Fisheries, 4600 Rickenbacker Cswy., Miami, FL 33149, USA

¹University of Western Sydney, School of Natural Sciences, Hawkesbury H4, Locked Bag 1797, Penrith South DC, NSW 1797, Australia (pm.ross@uws.edu.au)

²NSW Department of Primary Industries, Port Stephens Fisheries Centre, Taylors Beach, NSW 2316, Australia

LATITUDINAL VARIATION IN CALCIFICATION: VULNERABILITY OF ANTARCTIC BENTHIC CALCIFIERS TO OCEAN ACIDIFICATION [C]

Watson, Sue-Ann¹, Paul A. Tyler¹ and Lloyd S. Peck²

Heavily calcified marine animals such as molluscs, brachiopods and echinoderms have essential roles in ocean ecosystems and contribute to a large proportion of the total biomass in Antarctic benthic communities. How will these heavily calcified animals cope in a high-CO₂ world? Ocean acidification caused by anthropogenic CO₂ emissions will affect the ability of these calcareous organisms to build their shells and skeletons, particularly in the polar oceans where even modest CO₂ emission scenarios predict aragonite structures will enter a dissolution state by 2100. In order to assess the likely effects of ocean acidification on these calcareous invertebrates, we present data from polar, temperate and tropical latitudes on shell and skeleton size, calcium carbonate crystal form and elemental composition. Metabolic data were used to calculate the cost of shell production and wavelength dispersive spectroscopy was used to determine variation in shell composition between latitudes. Results show Antarctic snails, brachiopods and sea urchins have thinner shells and less shell per unit body mass than closely related temperate and tropical species. Scanning electron microscope imaging of crystal type shows shells of Antarctic bivalves and gastropods are composed predominately of aragonite. Having a smaller, more soluble shell raises particular concern for Antarctic molluscs as a reduction in shell deposition could increase their vulnerability to predation.

IMPACTS OF OCEAN ACIDIFICATION ON CALCAREOUS ZOOPLANKTON [I] Fabry, Victoria J.¹, Stephen Eggins², Bärbel Hönisch³, Fabien Lombard⁴, and Howard J. Spero⁵

Oceanic uptake of anthropogenic CO₂ is changing the seawater chemistry of the world's oceans towards reduced pH conditions, with profound consequences for many marine organisms. Marine zooplankton are fundamental to food webs, carbon dynamics, nutrient recycling and other ecosystem processes, yet little information exists on the response of many groups of these heterotrophs to elevated PCO₂. Increased PCO₂ in seawater can impact marine zooplankton both via decreased calcium carbonate saturation state, which may affect rates of shell secretion and dissolution, and via physiological processes involving acid-base reactions. Evidence suggests that two important groups of calcareous heterotrophs — the aragonitic pteropod molluses and calcitic planktonic foraminifers are sensitive to elevated PCO₂/reduced pH conditions. Only three species of these groups have been examined, however, and no direct measurements of calcification rates as a function of calcium carbonate saturation state are available. Shelled pteropods in the Southern Ocean are thought to be particularly at risk from ocean acidification because surface waters in that region may become persistently undersaturated with respect to aragonite as early as 2050. This presentation will focus on planktonic foraminifers and euthecosomatous pteropods. The available data will be summarized and new observations will be discussed to assess the potential impacts of present and projected changes in ocean carbonate chemistry on foraminifer and pteropod calcification rates. Critical knowledge gaps will be identified and directions for future research will be suggested.

School of Ocean and Earth Science, University of Southampton, National Oceanography Centre Southampton, European Way, Southampton, SO14 3ZH, United Kingdom (suwa@noc.soton.ac.uk)
 British Antarctic Survey, High Cross, Madingley Road, Cambridge, CB3 0ET, U.K.

¹Department of Biological Sciences, California State University San Marcos, San Marcos, CA 92096-0001, USA (fabry@csusm.edu)

²Research School of Earth Sciences, The Australian National University, Canberra 0200, Australia

³ Lamont-Doherty Earth Observatory, Department of Earth and Environmental Sciences, Columbia University, Palisades, NY 10964, USA

⁴DTU aqua, Jaegersborg Alle 1, DK-2920 Charlottenlund, Denmark

⁵Department of Geology, University of California Davis, Davis, CA 95616, USA

INTERANNUAL VARIABILITY OF PTEROPOD SHELL WEIGHTS IN THE HIGH-CO₂ SOUTHERN OCEAN [C]

 $\frac{Roberts,\ Donna}{Stephen\ G.\ Bray^{1}},\ William\ R.\ Howard^{1},\ Andrew\ D.\ Moy^{1},\ Jason\ L.\ Roberts^{1,2},\ Thomas\ W.\ Trull^{1,3,4},$ $Stephen\ G.\ Bray^{1}\ and\ Russell\ R.\ Hopcroft^{5}$

Anthropogenic CO₂ added to the atmosphere during and since the industrial revolution is entering the global ocean. Addition of extra (anthropogenic) CO₂ to the ocean changes the carbonate chemistry and the pH of the surface ocean, decreasing the concentration of carbonate ions and increasing ocean acidity. The ecological effects of changing ocean carbonate chemistry are uncertain, but are believed to include reductions in the calcification rates of marine organisms such as shelled pteropods.

The Southern Ocean contains a disproportionate amount of the oceanic inventory of anthropogenic CO₂, making it a biogeochemical harbinger for the impacts of ocean acidification, which may spread throughout the global ocean and might be expected to contain some of the first signals of impacts of increasing atmospheric CO₂.

Through an *in situ* sustained monitoring sediment trap deployed at 47°S in the deep (2000 m) Southern Ocean we've measured a significant drop in shell weight of a common cold-water Southern Ocean pteropod, *Limacina helicina antarctica* forma *antarctica*, of $1.17 \pm 0.47 \,\mu g \, yr^{-1}$ since 1997/98. With increasing levels of CO₂ entering the Southern Ocean the potential for further shell weight loss for this, and other, cold-water marine calcifiers could result in the change of distribution or complete removal of keystone species from the Southern Ocean ecosystem.

² Australian Antarctic Division, Channel Highway, Kingston, Tasmania, 7050, Australia

OCEAN CARBON SYSTEM: PAST & PRESENT

CONTROLS ON EVOLUTION OF OCEAN CARBONATE CHEMISTRY OVER THE PAST BILLION YEARS [I]

Caldeira, Ken

On time-scales that are long relative to the residence time, whatever materials enter the oceans (e.g., CO₂ from volcanoes etc, Ca²⁺ in rivers from rock weathering) must be balanced by materials deposited to the sediments. On long timer scales (> 10⁴ yr), the carbonate mineral saturation states of the ocean adjust to supply CaCO₃ to the sediments at the rate needed to balance inputs. The future high CO₂ ocean will be less saturated than today's whereas those of the extended high CO₂ (and high weathering rate) periods of the ancient past were likely more saturated. I will describe results of various methods to estimate past ocean chemistry, including both modeling and more observationally based approaches. Qualitative trends in paleo-ocean chemistry are clear, but quantitatively uncertain. The trend through evolutionary time is towards more efficient calcifiers, which would likely result in decreasing ocean saturation states. Thus, the ocean of the distant past it is likely to have been more saturated with respect to carbonate minerals than was the modern ocean. If current carbon dioxide emissions trends are not reversed, surface ocean pH will be lower than it has been in at least tens of millions of years and perhaps hundreds of millions of years. If these trends are not reversed, later this century ocean carbonate mineral saturation states could be lower than at any time in the past billion years with the exception of rare catastrophic events in Earth history such as the extinction events 65 and 250 myr ago.

¹ Antarctic Climate & Ecosystems Cooperative Research Centre, Hobart, Tasmania, 7001, Australia (D.Roberts@acecrc.org.au)

³ Commonwealth Scientific and Industrial Research Organisation (CSIRO) Marine and Atmospheric Research, Castray Esplanade, Hobart, Tasmania, 7000, Australia

⁴ Institute of Antarctic & Southern Ocean Studies, University of Tasmania, Hobart, Tasmania, 7001, Australia

⁵ Institute of Marine Science, University of Alaska Fairbanks, Fairbanks, Alaska 99775-7220, USA

Carnegie Institution, Department of Global Ecology, 260 Panama St, Stanford, CA 94305 USA (kcaldeira@stanford.edu)

BORON ISOTOPE EVIDENCE OF OCEAN ACIDIFICATION IN THE NEOPROTEROZOIC [C]

Kasemann, Simone A.¹, Anthony R. Prave² and Anthony E. Fallick³

The Neoproterozoic is a period of exceptional Earth System change marked by extreme fluctuations from icehouse to greenhouse climatic conditions. These severe environmental changes are preserved in the geological record of warm water carbonates sharply overlying glaciogenic strata. In concert with the environmental changes, the carbonate rocks display large amplitude fluctuations in their stable isotopic composition. These fluctuations are stratigraphically systematic, occur in many sections worldwide and are interpreted as being globally significant. Thus, the Neoproterozoic carbonates provide a unique geological and isotopic archive to improve our understanding of major non-anthropogenically influenced changes in Earth System behaviour.

The most intriguing isotopic signal is a characteristic negative C-isotopic excursion (down to -6 %). This signal has been used to assess atmospheric pCO_2 , organic productivity and carbon cycling during the extreme Earth System changes. Because of the relationship between carbonate $\delta^{13}C$, pCO_2 , ocean acidification and B isotopic composition of seawater, we analysed Neoproterozoic carbonates to obtain their B-isotopic signature and reconstruct ocean pH variation. We collected a B-C-O isotopic dataset on postglacial cap carbonates from two discrete glacial intervals in Namibia (~750-630 Ma). Systematic B-isotope excursions (up to 9‰) in the postglacial carbonates appear to be associated with fluctuations in atmospheric pCO_2 and changing weathering rates and can be interpreted as directly reflecting periods of acidification of the surface ocean. These data provide new insights into the nature of the initiation and termination of Neoproterozoic glaciations and into the B-isotopic composition of ancient oceans.

REDUCED CALCIFICATION IN MODERN SOUTHERN OCEAN PLANKTONIC FORAMINIFERA [C]

Moy, Andrew D.^{1, 2}, William R. Howard¹, Stephen Bray¹ and Tom Trull^{1, 3, 4}

Laboratory experiments suggest that decreased ocean carbonate-ion concentrations due to anthropogenic CO_2 will result in reduced calcification rates in marine calcareous organisms. Here we use sediment trap and Holocene sediment evidence to show reduced calcification in Southern Ocean planktonic foraminifera since the industrial revolution. Our sediment trap results provide the first field observations for reduced calcification (inferred from planktonic foraminiferal shell weights) in Southern Ocean carbonate organisms and indicate there has been a 30-34% reduction in calcification (in the planktonic foraminifer *Globigerina bulloides*), consistent with the impact due to altered ocean chemistry resulting from anthropogenic CO_2 storage. Down-core shell weight variations show reduced calcification during glacial-interglacial CO_2 increases that are similar to the post-industrial increase. Planktonic foraminifera contribute $\sim 23-50$ % of the total open-marine carbonate flux in the modern ocean, thus a reduction in the calcification of planktonic foraminifera may imply a corresponding decrease in calcium carbonate export from the surface ocean.

¹ School of Geosciences, Grant Institute of Earth Science, University of Edinburgh, Edinburgh EH9 3JW, U.K. (Simone.Kasemann@ed.ac.uk)

² School of Geography & Geosciences, University of St Andrews, St Andrews, Fife KY16 9AL, U.K.

³ Scottish Universities Environmental Research Centre, Scottish Enterprise Technology Park, Glasgow G75 OQF, U.K.

¹ Antarctic Climate and Ecosystems Cooperative Research Centre, University of Tasmania, Hobart, Australia

CURRENT RATES OF CHANGE IN PH AND CALCIUM CARBONATE SATURATION IN THE HIGH LATITUDE NORTH ATLANTIC OCEAN [C]

Olafsson, Jon¹, Solveig R. Olafsdottir¹, Alice Benoit-Cattin¹, Magnus Danielsen¹ and Taro Takahashi²

The rates of change in pH and calcium carbonate saturation are evaluated from observations at two time series stations in the vicinity of Iceland over the period 1983-2006. One station is in the northern Irminger Sea with relatively warm and saline Modified North Atlantic Water derived from the North Atlantic Current. The other is in the Iceland Sea where cold Arctic Intermediate Water usually predominates but the Polar Water influence in the surface layers is variable. These are source regions of the North Atlantic Deep Water. The pH and \square values were calculated from TCO2 and pCO2 observations.

Seasonal surface water observations show large variations on account of biological productivity, respiration and mixing. For pH the observed seasonal range in the Irminger Sea is 0.17 and for the Iceland Sea it is 0.23 pH units.

Winter observations, when biological activity is at seasonal minimum, show that the local rates of pH change are for the Irminger Sea -0.0012 yr⁻¹ and -0.0025 yr⁻¹ for the Iceland Sea. We show that the influence of hydrographic variability, e.g. temperature, on pH and calcium carbonate saturation must also be considered. The rates of change in calcium carbonate saturation decrease with depth in the Iceland Sea. However, at about 1800 m depth there the aragonite saturation is in transition to a state of undersaturation. Considering the hypsographic characteristics of the Iceland Sea, this transition may influence large areas of the sea floor.

LOW WINTER CACO₃ SATURATION IN THE BALTIC SEA AND CONSEQUENCES FOR CALCIFIERS [C]

Tyrrell, Toby¹ and Bernd Schneider²

We should attempt to understand the consequences of ocean acidification through a variety of techniques: not only through laboratory and mesocosm experiments but also through field observations of living communities in low pH (low saturation state) environments. Such observations are likely to be most useful when made in extreme environments that resemble in at least some regard the acidification-related changes predicted for the future.

Carbonate chemistry measurements in the Baltic Sea show that in winter it experiences the lowest surface saturation states yet observed for any ocean or sea region; the central Baltic Sea is undersaturated with respect to aragonite and borderline undersaturated ($\Omega \approx 1$) with respect to calcite. CaCO3 saturation state is even lower in the northern parts of the Baltic Sea. We will discuss the success or failure of some calcifying organisms (with particular emphasis on coccolithophores) to cope with this winter undersaturation, and the implications for their likely fate in a future more acidic ocean.

² Department of the Environment, Water, Heritage and the Arts, Australian Antarctic Division, Kingston 7050, Australia

³ University of Tasmania, Hobart, Tasmania

⁴ CSIRO Marine and Atmospheric Research, Hobart, Tasmania, Australia

¹ Marine Research Institute, Skulagata 4, 121 Reykjavik, Iceland. jon @ hafro.is

² Lamont-Doherty Earth Observatory, Palisades, NY 10964, USA

¹ National Oceanography Centre, Southampton University, European Way, Southampton SO14 3ZH, U.K. (tt@noc.soton.ac.uk)

² Institut für Ostseeforschung Warnemünde, Seestrasse 15, D-18119 Rostock, Germany.

EFFECTS OF OCEAN ACIDIFICATION ON NUTRIENT AND METAL SPECIATION

EFFECTS OF CHANGES IN CARBONATE CHEMISTRY ON NUTRIENT AND METAL SPECIATION [I]

De Baar, Hein^{1,2}, Loes Gerringa² and Charles-Edouard Thuroczy²

Ocean ecosystems tend to be limited due to limited availability for biological uptake of major nutrients (N, P, Si) and trace nutrients (Fe, Zn, Co, Mn, Cu, Cd) essential for life. Until recently the essential major element carbon (C) was deemed to be available in excess, but since about one decade we realize its chemical speciation (HCO₃-, CO₃²⁻) does affect uptake and growth by phytoplankton. Similarly the major nutrients P and Si exist as weak acids in seawater, with their chemical speciation depending on the ambient pH. Hence the ongoing acidification of surface ocean waters due to invasion of anthropogenic CO₂ will affect their chemical speciation, and this in turn may, or may not, affect their uptake by phytoplankton.

The chemical speciation hence biological availability of all trace nutrients (Fe, Zn, Co, Mn, Cu, Cd) is affected by the ongoing change in ocean carbonate chemistry, both directly in their speciation with HCO₃⁻ and CO₃² but even more indirectly due to the acidification, the latter also affecting the oxidation/reduction chemistry of crucial elements Fe, Co and Mn. Iron (Fe) is the dominant limiting nutrient in circa 40% of the world oceans, and therefore is of most interest, but happens to have a very complicated and dynamic chemistry in seawater, including its strong organic complexation. Moreover there is the formation and dissolution of Fe-colloids which in themselves are deemed unavailable for uptake by plankton. Ocean acidification affects both the organic complexation and the colloid state of Fe in seawater.

OCEAN ACIDIFICATION EFFECTS ON IRON SPECIATION IN SEAWATER [C]

Breitbarth, Eike^{1, 2}, Richard J. Bellerby², Craig C. Neill², Murat V. Ardelan³, Michael Meyerhöfer⁴, Eckart Zöllner⁴, Peter L. Croot⁴, and Ulf Riebesell⁴

The changing pH of seawater may influence the speciation of bio-relevant trace metals in the future oceans. We investigated the consequences of ocean acidification in coastal mesocosm CO_2 perturbation experiments, studying natural phytoplankton blooms under atmospheric CO_2 scenarios of 350, 700, and 1050 μ atm pCO_2 . Our experiments show significant increases in dissolved iron (<0.2 μ m) concentrations for the high CO_2 treatments during and after the blooms. Furthermore, the future ocean scenarios revealed higher Fe^{2+} values than the lower CO_2 treatments. Our findings suggest that biological feedback mechanisms in the future scenarios result in increased organic complexation of iron and thus increase the dissolved fraction of iron. Iron chelates are more photolabile at lower pH and together with slowed Fe^{2+} oxidation rates result in the increased Fe^{2+} concentrations. These trends may lead to higher iron bioavailability and could thus provide a negative feedback mechanism to the rising atmospheric CO_2 by fuelling primary production in the future ocean. This may be particularly relevant, if the mechanisms observed here also apply to currently iron limited open ocean high nutrient low chlorophyll (HNLC) regions.

¹ Ocean Ecosystems, Centre for Ecological and Evolutionary Studies, University of Groningen, PO Box 14, 9750 AA Haren, The Netherlands

² Royal Netherlands Institute for Sea Research, PO Box 59, 1790 AB Den Burg, The Netherlands (debaar@nioz.nl)

Department of Chemistry, University of Gothenburg, Sweden (eike@chem.gu.se)

² Bjerknes Centre for Climate Research, University of Bergen, Norway

³ Norwegian University of Science and Technology, Trondheim, Norway

⁴ Leibniz Institute of Marine Sciences, IFM-GEOMAR, Kiel, Germany

DAY 2: TUESDAY, 7 OCTOBER

MECHANISMS OF CALCIFICATION

BIOMINERALIZATION MECHANISMS IN MARINE CALCIFYERS IN VIEW OF OCEAN ACIDIFICATION [I]

Erez, Jonathan, Shmuel Bentov, Alon Braun, Mor Grinstein, Jack Silverman, Kenneth Schneider, and Boaz Lazar

Biomineralization of CaCO₃ is major process in the global carbon cycle. The major calcifyers in the ocean are Coccolithophores, foraminifera, corals and mollusks. Atmospheric CO₂ increase lowers the pH of the surface ocean and consequently the carbonate ion concentration is decreasing. While reports are conflicting with respect to coccolithophores, experiments and observations on corals and foraminifera show dramatic decrease in the rates of calcification with lower carbonate ion concentrations. This is also true for community calcification of an entire coral reef ecosystem. The sensitivity of corals and foraminifera to ocean acidification can be readily explained because both groups bring seawater to the site of biomineralization. Using confocal microscopy we found that in foraminifera, seawater is supplied to the biomineralization site by vacuolization. In corals we can demonstrate that cell-impermeable fluorescent probes (Calcein and FITC-dextran) added to the seawater are incorporated into the growing coral skeleton, suggesting the presence of seawater at the calicoblastic space. While the exact mechanism for the seawater supply is unknown, it may be associated with paracellular pathways, or gaps between the tissue and the porous skeleton. In both groups the main modification that the organisms exert on the seawater in order to calcify is elevation of pH by roughly 1 pH unit relative to ambient levels. Ocean acidification increases the pH elevation gap as well as the buffer capacity of the seawater, thus even a small pH decrease of 0.2-0.3 pH units is strongly affecting the calcification rates. Implications for other groups will be discussed.

Institute of Earth Sciences, The Hebrew University of Jerusalem, Jerusalem 91904, Israel. (erez@vms.huji.ac.il)

EFFECT OF ACIDIFICATION ON CORAL CALCIFICATION: WORKING HYPOTHESES TOWARDS A PHYSIOLOGICAL MECHANISM [C]

Marubini, Francesca, Denis Allemand and Christine Ferrier-Pagès

In order to determine what are the factors involved in the seawater acidification-induced decrease of coral calcification, we assessed the effect of seawater acidification on the calcification and photosynthesis of the scleractinian tropical coral *Stylophora pistillata in different physico-chemical conditions*. Coral nubbins were incubated for 8 days at three different pH (7.6, 8.0 and 8.2) obtained by adding HCl or NaOH. For each pH, corals were maintained under bicarbonate-ambient (2 mM) and bicarbonate-enriched (4 mM) conditions. Bicarbonate-enriched condition was achieved by adding into the tanks 2 mM bicarbonate (HCO₃) using a peristaltic pump. Photosynthesis of *S. pistillata* was insensitive to changes in pH and pCO₂, but was increased in the tanks enriched with 2 mM HCO₃, suggesting carbon-limited conditions. Seawater acidification decreased coral calcification by ca. 0.1 mg CaCO₃ g⁻¹ d⁻¹ for a decrease of 0.1 pH units. As previously shown, decrease in calcification was correlated with a decrease in the availability of the CO₃²⁻ substrate, and thus a decrease in the saturation state, W. However, our results show that the decrease in coral calcification could also be attributed either to a decrease in extra-or intracellular pH or to a change in the buffering capacity of the medium, impairing supply of CO₃²⁻ from HCO₃.

Centre Scientifique de Monaco, Avenue Saint-Martin, MC-98000 Monaco, Principality of Monaco. Email: allemand@centrescientifique.mc

PREDICTION OF CARBON FIXATION DURING A BLOOM OF EMILIANIA HUXLEYI IS HIGHLY SENSITIVE TO ASSUMED RESPONSE TO SHIFT IN pCO_2 [C]

Bernard, O.¹, A.Sciandra² and S.Rabouille²

Large scale precipitation of calcium carbonate in the oceans by coccolithophorids is a phenomenon that plays an important role in carbon sequestration. However, there is a controversy on the effect of an increase in atmospheric CO₂ concentration on both calcification and photosynthesis of coccolithophorids. Indeed recent experiments, performed in conditions of nitrogen limitation, revealed that the associated fluxes may be slowed down, while other authors claim the reverse response. We have designed models to account for various scenarii of calcification and photosynthesis regulation in chemostat cultures of Emiliania huxleyi, based on different hypotheses of regulation mechanism. These models, which are kept at a general and generic level, consider that either carbon dioxide, bicarbonate, carbonate or pH is the regulating factor. These models are calibrated to predict the same carbon fluxes in nowadays pCO₂, but they turn out to respond differently to an increase of CO₂ concentration. Thus, we simulated a bloom of Emiliania huxleyi using the 4 considered regulation scenarii. For high biomass concentration, the coccolithophorids can significantly affect the inorganic carbon and the pH in their environment, thus leading to a feedback in their growth rate which is, depending on the model, positive or negative. It results that the prediction of the carbon fixed during the bloom varies by a factor 2, depending on the assumed regulating mechanism hypothesized for growth and calcification.

¹INRIA-COMORE, BP93 Sophia-Antipolis Cedex, France (olivier.bernard@inria.fr) ²LOV, UMR 7093, Station Zoologique, B.P. 28 06234, Villefranche-sur-mer, France

PHYSIOLOGICAL EFFECTS: FROM MICROBES TO FISH

PHYSIOLOGICAL MECHANISMS LINKING CLIMATE TO ECOSYSTEM CHANGE: EFFECTS OF OCEAN ACIDIFICATION ON MARINE ANIMALS IN TIMES OF OCEAN WARMING [I]

Pörtner, Hans O.

Climate change causes ocean warming and acidification on global scales. In contrast to well established effects of warming on marine ecosystems, specific effects of rising CO₂ are expected but evidence is still equivocal. However, future scenarios indicate a threatening of marine life through the specific or synergistic effects of rising CO₂ levels, warming and more frequent hypoxia events. Beyond empirical observations, development of a cause and effect understanding and of realistic scenarios is required for a secure projection of ecosystem effects. Such understanding builds on the identification of key physiological mechanisms and their responses to progressive acidification, warming and hypoxia. In changing oceans, these are physiological characters which define species performance, including their capacity to interact, e.g. in food webs. Many current ecosystem changes likely occur when ambient temperature drifts beyond species specific thermal tolerance windows and causes a shift in phenology. Specific sensitivity to elevated CO₂ levels may involve a key role for acid-base regulation, with low capacities found in lower marine invertebrates. Extracellular acid-base status feeds back on the costs of ion and acid-base regulation, on processes involved in growth, on calcification, neural functions, blood gas transport or behavioural capacities. The capacities of metabolic pathways shift to new equilibria. Present evidence indicates elevated sensitivity to elevated CO₂ levels towards the edges of thermal windows. The key consequence may be a narrowing of thermal tolerance windows and associated ranges of geographical distribution, of the scope for performance at ecosystem level and thus, an exacerbation of warming effects on marine ecosystems.

Alfred-Wegener-Institute for Polar and Marine Research, Marine Animal Physiology, Bremerhaven, D-27515, Germany (hans.poertner@awi.de)

IMPACTS OF OCEAN CHANGE ON PRIMARY PRODUCERS [I]

Riebesell, Ulf

Rising atmospheric CO₂ and the resulting climate change cause the ocean to undergo two major alterations: sea surface warming and ocean acidification/carbonation. While there is strong evidence that both of these changes are in progress at an ocean-wide scale, little is known about their individual and combined effects on the marine biota. Hence, our ability to forecast the resulting impacts on ocean productivity, elemental cycling and air/sea exchange is rudimentary and model scenarios of future ocean changes are mostly based on simplifying concepts of possible ecosystem and biogeochemical alterations.

Despite much uncertainty about the impacts on marine biota, there is growing evidence that there will be both winners and losers of ocean carbonation (increasing pCO₂) and acidification (decreasing pH). Calcareous organisms will for the most part be on the losing side as increasing seawater acidification incurs a greater metabolic energy requirement to precipitate calcium carbonate. Some photoautotrophic groups are likely to be on the winning side as increasing ocean carbonation makes it energetically less expensive to obtain the CO₂ required for photosynthesis. In addition, sea surface warming effects marine biota both directly, due to the strong temperature sensitivity of most biological processes, and indirectly through changes in surface layer stratification, overturning circulation, and the resulting effects on nutrient supply and light intensities. This presentation will review the observed biotic responses in pelagic systems and assess their possible impacts on ocean productivity, biogeochemical cycling and their feedback potential to the climate system.

CO2 LEAKAGE IN THE DEEP OCEAN AND ITS EFFECT ON BIOTA AND BIOGEOCHEMISTRY – LESSONS FROM NATURAL ANALOGUES ON ${\rm CO_2}$ DISPOSAL IN THE OCEAN [I]

Boetius, Antje^{1,2}, Dirk de Beer¹, Judith Ufkes¹, Matthias Haeckel³, Fumio Inagaki⁴, Koichi Nakamura⁵, Gregor Rehder⁶

One of the future strategies to deal with excess CO₂ in the atmosphere is its disposal and storage in the ocean. The environmental risks of this mitigation strategy are not well constrained. Critical questions to any disposal scenario are as to the tipping points at which endemic species are affected by increasing pCO₂ and decreasing pH, and as to the threshold for CO₂ leakage from the seafloor to prevent negative effects on biodiversity and ecosystem services. During a recent expedition with FS SONNE and ROV QUEST to the Okinawa Trough we have investigated the potential effects of natural CO₂ leakage on deep-water benthic ecosystems at 1350 m water depth. At the Yonaguni Knoll vent system, volcanic CO₂ emissions lower the pH of the bottom water down to 7.2, with dramatic effects on megafauna distribution and composition. Furthermore, subsurface accumulations of liquid CO₂ cause a decrease in pH down to pH 4.5 in the ocean seafloor, altering considerably biogeochemical processes in the sediments and affecting microbial communities. This presentation discusses direct and indirect effects of CO₂ accumulation and leakage on deep-water ecosystems and relevant biogeochemical processes.

- 1 Max Planck Institute for Marine Microbiology, Celsiusstr.1, 28359 Bremen, Germany
- 2 Jacobs University Bremen, 28759 Bremen, Germany
- 3 Leibniz-Institute for Marine Sciences IFM-GEOMAR, Wischhofstr. 1-3, 24148 Kiel, Germany
- 4 Kochi Institute for Core Sample Research, JAMSTEC, Monobe B200, Nankoku, Kochi 783-8502, Japan
- 5 National Institute of Advanced Industrial Science and Technology (AIST), Institute of Geology and Geoinformation, 7, 1-1-1- Higashi, Tsukuba, Ibaraki 305-8567, Japan
- 6 Baltic Research Institute, Seestraße 15, 18119 Rostock-Warnemünde, Germany

¹ Leibniz Institute of Marine Sciences, IFM-GEOMAR, Kiel, Germany (uriebesell@ifm-geomar.de)

EFFECTS OF HYPERCAPNIC ACIDIFICATION OF SEAWATER ON THE BIOLOGY OF NON-CALCIFYING MARINE ORGANISMS [C]

Thuesen Erik V.¹ and Brad A. Seibel²

Human activity is quickly increasing the carbon dioxide concentration (PCO₂) in the Earth's atmosphere, resulting not only in rapid climate change but an increase in the PCO₂ content of ocean waters above normal levels, a condition know as hypercapnia that leads to a decrease in the pH of seawater. The acidification of ocean waters due to hypercapnia can have extensive effects on marine life. For example, snails and bivalves may not be able to maintain their calcified shells and calcium carbonate reefs may dissolve due to the equilibrium shifts in the carbonate in seawater. For noncalcifying organisms, a whole other suite of problems may occur. This presentation reviews the potential effects of hypercapnic acidification on biological processes not related to calcification. It has been proposed that animals with lower metabolic potentials will be the most challenged by declining ocean pH because they lack adequate buffering systems and ion exchange mechanisms to cope with intracellular acidosis promoted by extracellular hypercapnia. Other examples include the following: increased metabolic cost to maintain intracellular pH, lower protein synthesis and decreased growth rates, proper function of transmembrane proteins, the production and function of mucus, cell-cell signaling. These and other biological impacts are reviewed and mechanistic models are presented in order to begin to prioritize productive areas of research into the effects of ocean acidification on noncalcifying marine organisms.

PREDICTING THE IMPACT OF OCEAN ACIDIFICATION ON BENTHIC BIODIVERSITY: WHAT CAN ANIMAL PHYSIOLOGY TELL US? [C]

Widdicombe, Stephen¹ and John I. Spicer²

For the past 200 years, the oceans have been absorbing carbon dioxide at an unprecidented rate. It is now evident that this ongoing process has already significantly altered seawater carbon chemistry at a global scale and will continue to do so for hundreds of years to come; a phenomenon termed "ocean acidification". The challenge currently facing scientists is to predict the long term implications of ocean acidification for the diversity of marine organisms and for the ecosystem functions this diversity sustains. This challenge is all the more difficult considering that empirical data which specifically address the impact of ocean acidification on marine biodiversity are currently lacking. In the face of growing political and public pressure to provide answers, what predictions can be made and how reliable are the assumptions on which those predictions depend? Here we review the extent to which the few existing data, and understanding gained from previous physiological studies, can be used to make predictions for marine biodiversity. In doing so we also scrutinise some established paradigms concerning the impact of hypercapnia, resulting in seawater acidification, on marine organisms.

¹ Evergreen State College, Laboratory I, Olympia, WA 98505, USA (thuesene@evergreen.edu)

² Department of Biological Sciences, University of Rhode Island, 100 Flagg Road, Kingston, RI 02881, USA.

¹ Plymouth Marine Laboratory, Prospect Place, West Hoe, Plymouth, PL1 3DH, U.K. (swi@pml.ac.uk)

² Marine Biology and Ecology Research Centre, School of Biological Sciences, University of Plymouth, Plymouth PL4 8AA, U.K.

FISHERIES, FOOD WEBS, AND ECOSYSTEM IMPACTS (PICES-ICES SESSION)

CONSEQUENCES OF OCEAN ACIDIFICATION FOR FISHERIES [I]

Fosså, Jan Helge¹, Richard Bellerby² and Tore Jakobsen¹

The distribution of major fisheries is compared with model predictions of pH change in different regions of the world ocean to identify fisheries in ecosystems that may be affected by a change in ocean chemistry and associated biogeochemical and ecological change. The potential impact of ocean acidification on marine fish will be assessed by reviewing how higher seawater CO₂, and associated changes to the marine carbonate system, can affect, both directly and indirectly, the reproduction, growth and mortality of fish populations. Case studies of collapsed fish stocks will be presented to illustrate how a combined effect of environmental change and high fishing pressure can be detrimental to fish stocks and the fisheries that depend upon them.

There has been no direct research on the consequences of ocean acidification for fisheries. There is, however, growing evidence of effects of ocean acidification on important processes within marine ecosystem functioning and on key organisms in the marine food web. On this basis, potential impacts from ocean acidification on ecosystem structure and function related to the position of fish in the food webs will be discussed for a range of ecosystems, e.g. the Norwegian Sea, the Benguela upwelling system, and tropical coral reefs.

The main conclusion is that the threat from ocean acidification urgently calls for a fisheries management that aims at maintaining strong and robust fish stocks with minimum loss of genetic variation in order to secure maximum potential for adaptation to the expected and possibly irreversible changes in the environment.

² Bierknes Center for Climate Research, Allégaten 55, N-5007 Bergen, Norway

AN OCEAN ACIDIFICATION SIMULATION EXPERIMENT WITH BENTHIC ANIMALS USING A PRECISE PCO₂ CONTROL SYSTEM [I]

<u>Nojiri, Yukihiro</u>¹, Yoshihisa Shirayama², Hideshi Kimoto³, Takeshi Egashira³ and Katsumoto Kinoshita³

Japan started a research program entitled as 'Experimental study of ocean acidification impact on benthic calcifiers' funded for 2008-2010 by Ministry of Environment. In this program, precise CO₂ controlling system for culturing benthic calcifiers, such as sea urchin, shellfish and coral, will be operated to experiment the impact of CO₂ increase on the coastal benthic animals. Preliminary study and numerical consideration demonstrate difficulties in controlling seawater pCO₂ using a regular aquarium tank and gas bubbling. It is because of insufficient equilibration efficiency with several tens centimeters of bubbling height in the tank. In sufficient equilibration is especially enhanced when aquarium is maintained by overflowing water. Carrying CO₂ away by the overflowing water gives insufficient equilibration and then it needs very high equilibration sufficiency of bubbling air. In case the experiment is done without monitoring pCO₂ of the aquarium tank may give an artifact. We developed a new type of air-seawater equilibrator by combination of a 'counter current dissolution water tower (CCDWT)' and a 'measurement water tower (MWT)'. The system can produce seawater with target pCO₂ and record pCO₂ by a LICOR gas analyzer. Seawater having target pCO₂ can be supplied to an aquarium tank at around 20L/hour rate, which enables the turnover time of several hours for regular size tank. The system can simulate diurnal cycle of pCO₂, generally occurs in coastal area. Our culture experiment has been stared to research long term and low level exposure of CO₂ for benthic animal species in Japanese coastal environment.

¹ Institute of Marine Research, P.O. Box 1870 Nordnes, N-5817 Bergen, Norway (jhf@imr.no)

¹ Natl. Inst. Environ. Stud., Tsukuba, Ibaraki 305-8506, Japan (nojiri@nies.go.jp)

² Seto Mar. Biol. Lab., Kyoto Univ., Shirahama, Wakayama, 649-2211, Japan

NATURAL CO2 VENTS REVEAL ECOLOGICAL TIPPING POINTS DUE TO OCEAN **ACIDIFICATION [C]**

Hall-Spencer, Jason M. and Riccardo Rodolfo-Metalpa

Our understanding of how increased ocean acidity may affect marine ecosystems is currently severely hampered, since almost all studies have been in vitro, short-term, rapid perturbation experiments on isolated elements of the ecosystem. Here we show the effects of acidification on benthic ecosystems at shallow coastal sites where volcanic CO₂ vents lower the pH of the water column. Along gradients of normal pH (8.1-8.2) to lowered pH (mean 7.8-7.9), typical rocky shore communities with abundant calcareous organisms shifted dramatically to communities lacking scleractinian corals with significant reductions in sea urchin and coralline algal abundance. This is the first ecosystem-scale validation of predictions that these important groups of organisms are susceptible to elevated levels of pCO₂ and offers insights into community shifts to be expected in 2100 and beyond. Seagrass production peaked in an area at mean pH 7.6 (1827 µatm pCO₂) where coralline algae were eradicated and gastropod shells were dissolving due to periods of carbonate sub-saturation. Volcanic CO₂ vent sites reveal a suite of marine organisms that are resilient to naturally high levels of p CO₂, which would be difficult to predict using modelling and mesocosm approaches, and that ocean acidification may benefit highly invasive alien algal species. Our results provide the first in situ insights into how shallow water marine communities might change when susceptible organisms are removed due to ocean acidification.

Marine Institute, Biological Sciences, University of Plymouth, Plymouth PL4 8AA, U.K.

SALMON PHISHING IN THE NORTHEAST PACIFIC; AN ARCHAEOLOGICAL DIG IN THE NORTH PACIFIC SURVEY DATA (1956-1964) [C] McKinnell, Skip, Kerim Aydin, Jim Christian³, Nancy Davis, and David Mackas

Pacific salmon eat pteropods; yet until the threat of a shoaling aragonite saturation level in the North Pacific emerged, scant attention was paid to where, when, in what quantities, and with what preference. This presentation provides a synoptic view of the distributions of pH, pteropods, and Pacific salmon and their diets in the North Pacific during the only period in history when all were surveyed simultaneously. Following the establishment of the International North Pacific Fisheries Commission in the mid-1950s, comprehensive, multi-vessel surveys of the physical, chemical, and biological oceanography, and the biology of Pacific salmon were conducted in the Gulf of Alaska for nearly a decade. The data include hydrographic and chemical profiles from hundreds of bottle casts, zooplankton tows, and 10,000 stomach samples analyzed for diet. These data have been lying fallow for decades and their resurrection facilitates the creation of baseline conditions at the beginning of the "Keeling Curve" that can serve as a basis for comparisons with recent and future samples.

³ Kimoto Electric Co. Ltd., 3-1, Funahashi-cho, Tennoji-ku, Osaka 543-0024, Japan

¹ North Pacific Marine Science Organization, Sidney, Canada (mckinnell@pices.int).

² Alaska Fisheries Science Center, National Oceanic and Atmospheric Administration, Seattle, USA.

³ Fisheries and Oceans Canada, Institute of Ocean Sciences, Sidney, Canada

⁴ Fisheries Sciences, University of Washington, Seattle, USA

CO₂ DISPOSAL

EFFECTS OF CO₂ CAPTURE AND STORAGE ON OCEAN ACIDIFICATION [I] Haugan, Peter M.

The presently ongoing ocean acidification is the result of global scale indirect ocean storage of CO_2 via emissions to the atmosphere. The ocean uptake reduces the immediate greenhouse forcing on climate, but may increase stress on ocean ecosystems. In addition to this unintended indirect ocean storage, schemes for capture of CO_2 primarily from fossil fuel power plants and direct storage in the ocean or in geological reservoirs (CCS), have recently gained substantial support. The main aim of this paper is to provide a review of such schemes and discuss their possible future impact on local and large scale ocean acidification as well as global carbon cycling and climate.

Recent changes in global and regional conventions and regional and national legislation have opened up possibilities for direct storage of CO₂ in geological reservoirs on land or underneath the seabed on continental shelves. A range of projects are approaching implementation. If CO₂ leaks from offshore storage sites, local acidification will be strong. Due to the energy penalty associated with capture, the amount of CO₂ stored will be higher than that vented to the atmosphere in the case of no capture. Some greenhouse warming can be avoided by CCS compared to emissions, but the risk of leakage remains, with implications for future acidification and climate effects. Uncertainties about geological storage in hot rocks where CO₂ is buoyant, suggests that alternative deep ocean seafloor storage options may be reconsidered. In that case mechanisms for spreading into the water column become crucial for acidification effects.

Geophysical Institute, University of Bergen, Allegaten 70, N-5007 Bergen, Norway (Peter.Haugan@gfi.uib.no)

MODELLING OF CO₂ DISPERSION LEAKED FROM SEAFLOOR OFF JAPANESE COAST [C]

Yuki Kano¹, <u>Toru Sato</u>², Jun Kita³

A numerical study was conducted to predict the rise of pCO_2 in the ocean on a continental shelf by the leakage of CO_2 , which is originally stored in the aquifer under the seabed, in an extreme case, such as a large fault connects the CO_2 reservoir and the seabed accidentally by a big earthquake or other large diastrophism. The target space in this study was limited to the ocean above the seabed and does not include subsea underground. First, parameter studies were conducted in a small domain to see the impacts of CO_2 leakage and seawater conditions. The CO_2 takes a form of bubbles or droplets, depending on the depth of water, and their behaviour and dissolution were numerically simulated during the rise in constant background flows perpendicular to the fault. Also simulated was the advection-diffusion of dissolved CO_2 . Next, a more realistic case simulation was conducted with topography and tidal current at the offshore of a selected coast of Japan. As a result, it was suggested that the rises of CO_2 concentration were smaller than 500 ppm, even in such an extreme case with specific assumptions.

¹ Geological Survey and Applied Geoscience Laboratory, National Institute of Advanced Industrial Science and Technology (AIST), 1-1-1 Higashi, Tsukuba 305-8567, Japan

² Department of Ocean Technology, Policy, and Environment, University of Tokyo, 5-1-5 Kashiwanoha, Kashiwa 277-8563, Japan (sato-t@k.u-tokyo.ac.jp)

³ Demonstration Laboratory, Marine Ecology Research Institute, 4-7-17 Arahama, Kashiwazaki 945-0322, Japan

DAY 3: WEDNESDAY, 8 OCTOBER

ADAPTATION AND MICROEVOLUTION

A BRIEF HISTORY OF SKELETONS IN THE OCEANS [I]

Knoll, Andrew H.

Eukaryotes existed for more than a billion years before the biomineralized skeletons entered the fossil record. Consistent with phylogenetic evidence for the widespread use of silica by protists, the earliest mineralized elements are chromalveolate and rhizarian scales in 750-800 Ma rocks. Carbonate and silica skeletons occur as part of Ediacaran animal diversification, but a relatively late and taxonomically limited part. In contrast, carbonate, phosphatic and siliceous skeletons all diversified markedly as part of the greater Cambrian radiation of animal life. By the mid-Ordovician Period, animal and benthic algal skeletons had come to play much the same biomechanical, ecological and biogeochemical roles they do today. But the course of skeletal evolution never did run smooth. A remarkable decline in skeletal abundance and diversity occurred at the end of the Permian Period (252) Ma), when mass extinction devastated marine ecosystems. Patterns of extinction versus survival match predictions informed by physiological research on hypercapnia; rapid imposition high P_{CO2} at a time when deep ocean water masses already tended toward dysoxia may explain this greatest of all extinction events. Other mass extinctions show differing biological patterns, suggesting that Permo-Triassic scenarios cannot be applied to all such events. Episodic crashes of hypercalcifiers (including reef builders), however, may reflect transient but repeated intervals of high P_{CO2} and deep water anoxia in the world's oceans. Deep ocean oxygen levels have rarely been higher than they are at present. In this respect, at least, the present day challenges of ocean acidification depart from some deep time analogs.

Department of Organismic and Evolutionary Biology, Harvard University, Cambridge MA 02138, USA (aknoll@oeb.harvard.edu)

INFLUENCE OF HIGH CO_2 ON COCCOLITHOPHORES UNDER LONG-TERM CULTIVATION

Müller, Marius N., Kai G. Schulz, Peter Wiebe, and Ulf Riebesell

Studies on the sensitivity of coccolithophores to elevated CO₂ have generally relied on relatively short-term incubations over <20 and in some cases <5 generations. A critical issue in the assessment of ocean acidification impacts on marine biota is the ability of CO₂/pH sensitive organisms to adapt to changing environmental conditions. To approach this question we have exposed three coccolithophore species to gradually increasing CO₂ concentrations over more than 140 generations, namely *Emiliania* huxleyi (max. pCO₂ 1150 μatm), Coccolithus pelagicus braarudii (max. 930 μatm) and Calcidiscus leptoporus quadriperforatus (max. 740 µatm). All three species showed significant decreases in cellular division rates ranging from 10 to 30%, whereas organic matter production rates increased. Surprisingly, the increase of cellular organic matter was not reflected in an increase in cell diameter. In fact, C. pelagicus braarudii even shrank in size. Responses in cellular calcification rates were found to be species-specific as shown in previous studies. Whereas C. pelagicus braarudii reduced the calcification rate by 25 % and was covered with partly malformed coccoliths, an effect on E. huxleyi could not be detected due to high standard deviations. Additionally, we tested the effect of temperature in combination with high CO₂ on the performance of Emiliania huxleyi. Increased temperature amplified the decrease in cell division rate under high CO₂, but partly compensated the observed effect on calcification. These results are generally consistent with published data, but indicate a higher sensitivity of coccolithophores to ocean change than previously reported.

Leibniz Institute of Marine Sciences, 24105 Kiel, Germany (mnmueller@ifm-geomar.de)

NEW CONCERNS

IMPACT OF OCEAN ACIDIFICATION ON UNDERWATER SOUND: REDUCED LOW FREQUENCY ABSORPTION, INCREASED NOISE LEVELS, POTENTIALLY HIGHER STRESS FOR MARINE MAMMALS. [C]

Browning, David¹ and Peter M. Scheifele²

The principle absorption mechanism for low frequency, (<1,000 Hz), sound in the sea is a boron – boric acid reaction that is pH dependent. As a result, ocean acidification will reduce the low frequency sound absorption. To illustrate, the present difference between the average pH value in the ocean sound channel in the North Pacific Ocean (7.7) and that for the North Atlantic Ocean (8.0) results in the low frequency absorption being twice as large in the North Atlantic. Acidification will progress from the surface downward but since the dominant noise sources (shipping and wind wave generation) are also located at or near the surface, the impact on increasing noise levels should be starting also. As a result, marine mammals migrating at a shallow depth would be exposed to increasing noise levels which could potentially be a source of stress.

EXPERIMENTAL APPROACHESOF CARBONATE CHEMISTRY MANIPULATION IN CO_2 PERTUBATION STUDIES [C]

Schulz, Kai G. and Ulf Riebesell

Invasion of anthropogenic carbon dioxide (CO_2) into the surface ocean alters the seawater carbonate chemistry, increasing CO_2 and bicarbonate (HCO-3) at the expense of carbonate ion concentrations. This redistribution in the dissolved inorganic carbon (DIC) pool decreases pH and carbonate saturation state (Ω) . The latter, for instance, is thought to be a key variable determining calcium carbonate precipitation rates in many marine organisms such as coccolithophores, foraminifera and corals. The future survival of some marine ecosystems will rely on the success of calcifying organisms. Hence, more and more studies focus on the potential effects of changes in seawater carbonate chemistry on marine life. This requires accurate carbonate system manipulations and well controlled experimental setups. Here we describe and analyze the chemical consequences of increasing anthropogenic CO_2 in seawater. Furthermore we present several experimental approaches for carbonate chemistry manipulation, including acid/base, CO_2 and combined HCO-3 /acid additions, and compare their potential to simulate future ocean change. This will result in some fundamental guidelines for conducting CO_2 perturbation experiments.

Leibniz Institute for Marine Sciences (IFM-GEOMAR), D"usternbrookerWeg 20, 24105 Kiel, Germany (kschulz@ifm-geomar.de)

¹ Department of Physics, University of Rhode Island, 2 Lippitt Road, Kingston, RI 02881, USA (decibeldb@aol.com)

² Department Communications Sciences & Disorders, University of Cincinnati Medical Center, P.O. Box 670379, Cincinnati, OH 45267-0379, USA (scheifpr@ucmail.uc.edu)

BIOGEOCHEMICAL CONSEQUENCES AND FEEDBACKS TO THE EARTH SYSTEM

BIOGEOCHEMICAL CONSEQUENCES OF OCEAN ACIDIFICATION [I]

Bopp, Laurent

In the past 200 years, the ocean has absorbed half of the CO₂ emitted by human activities. This uptake has clearly slowed down anthropogenic climate change but it has led to changes in the chemical equilibrium of seawater and to a reduction of seawater pH by 0.1 since pre-industrial times. With increasing CO₂ emissions in this century, ocean surface pH could decrease by another 0.2-0.4 in 2100. The impacts of ocean acidification on marine biogeochemistry are still poorly understood. But we can ask the questions of the potential large-scale biogeochemical consequences and the potential to feedback on climate of such chemical changes?

Several lab experiments and field studies have shown that acidification would affect calcification, and hence the equilibrium between calcifying and non-calcifying species. I will review here the potential effects that these changes would produce on air-sea CO₂ exchanges and dimethylsulfide (DMS) emissions to the atmosphere.

In addition to calcification, acidification is also believed to potentially impact organic matter production, N-fixation, metal speciation.... I will review the potential effects of some of those modifications on the large-scale biogeochemistry of the ocean and on air-sea CO_2 and N_2O exchanges. In the last part, I will show how future climate change, mainly through changes of ocean circulation, could interact with acidification: dampening or amplifying acidification in the different regions of the world ocean but also amplifying or dampening some of the consequences of acidification

LSCE/IPSL, Orme des Merisiers, Bat 712, CE Saclay, F-91191 Gif sur Yvette, France. Laurent.Bopp@lsce.ipsl.fr

DISSOLUTION OF CaCO₃ IN SHALLOW WATER CARBONATE ENVIRONMENTS IN THE HIGH CO₂ WORLD OF THE ANTHROPOCENE [C]

Andersson, Andreas J.¹, Fred T. Mackenzie², Nicholas R. Bates¹, Ilsa B. Kuffner³, Abraham Lerman⁴, Paul L. Jokiel⁵, Ku'ulei S. Rogers⁵, and Adrian Tan².

Acidification of surface seawater owing to fossil fuels and land use change CO₂ emissions has raised serious concerns as to its consequences for marine organisms and ecosystems, especially those organisms producing shells, tests or skeletons of calcium carbonate. The effects of increasing seawater CO₂ and decreasing pH and carbonate saturation state on the rate and the ability of calcifying organisms to produce calcium carbonate have received significant attention. Much less attention has been given to the effects on carbonate mineral dissolution, although it is well known that dissolution of carbonate minerals will increase as a result of decreasing seawater carbonate saturation state. Numerical modeling simulations using the Shallow-water Ocean Carbonate Model (SOCM) demonstrate that the rate of dissolution of carbonate minerals in the global coastal ocean could exceed the rate of calcification in this region within the next century. Similarly, observations from a carbonate dominated environment in Bermuda exposed to elevated levels of CO₂ similar to projections for the future show carbonate dissolution rates sufficiently high to exceed the average rate of calcification on coral reefs within the present century. Furthermore, results from a mesocosm experiment subject to seawater conditions anticipated by the year 2100, show evidence of a net loss of carbonate material at the ecosystem scale, but also at the scale of individual organisms including coralline algae. These robust findings from three independent studies demonstrate that ocean acidification can push shallow water ocean environments and even some calcifying organisms to a state of net dissolution of carbonate material.

IMPACTS OF OCEAN ACIDIFICATION ON MARINE BIOGENIC TRACE GAS PRODUCTION [C]

Hopkins, Frances E.^{1, 2}, Suzanne M. Turner¹, Philip D. Nightingale², Michael Steinke³, and Peter S. Liss¹

The rapid and dramatic changes to ocean carbonate chemistry that will occur as result of ocean acidification are expected to have a detrimental impact on marine biological and biogeochemical processes. A potential consequence of this is that the sea-to-air flux of marine trace gases may be affected, which would impact on the oxidative capacity of the atmosphere, the production of particles in the marine boundary layer, air quality and climate. We have used three approaches to investigate whether trace gas production is affected by experimental changes in seawater pH: laboratory incubations of natural assemblages, a mesocosm experiment in Bergen, Norway, and a naturally-acidified coastal site in the Gulf of Naples, Italy. During the mesocosm experiment, the concentrations of a number of trace gases, including dimethyl sulphide (DMS) and volatile iodocarbon compounds, were significantly reduced under future atmospheric CO₂. We will present results from the recent fieldwork campaign in Italy, which will include an assessment of the suitability of the site as a model of the affects of future acidification on the pelagic community and associated biogeochemical processes.

FROM LABORATORY MANIPULATIONS TO EARTH SYSTEM MODELS: AN 'EPPLEY CURVE' FOR CALCIFICATION RATE? [C]

Ridgwell, Andy ¹, Daniela N. Schmidt ², and Carol Turley ³

The apparent incongruence between coccolithophore calcification responses observed across different experimental manipulations, particularly those involving *Emiliania huxleyi*, raises new challenges for modelers. This is because the global models used for predicting future fossil fuel CO₂ uptake by the ocean base their parameterizations for calcification closely on laboratory results. Predictions of such models will be unreliable if rooted in unrepresentative and/or poorly understood laboratory experiments.

To help address this, we have examined the specific details of available experimental manipulations. We find fundamental differences between laboratory culture studies, particularly in the strain (ecotype or likely even genotype) of E. huxleyi cultured. This, combined with the observations that: (1) experiments using other coccolithophore species have delineated the existence of a calcification 'optimum' in environmental conditions (pH), and (2) there is an unambiguous direction to the calcification- CO_2 response in mesocosm and shipboard incubations, lead us to propose a new formulation for describing ocean acidification impacts on planktonic marine carbonate production.

We suggest an equivalence can be drawn between species or even ecosystem integrated phytoplankton calcification rate as a function of pH (or saturation), and widely used descriptions of

¹Bermuda Institute of Ocean Sciences, 17 Biological Station, St George's GE01, Bermuda (andreas.andersson@bios.edu).

²Department of Oceanography, University of Hawaii, 1000 Pope Rd., Honolulu, HI 96822, USA.

³US Geological Survey, Florida Integrated Science Center, St Petersburg, FL 33701, USA.

⁴Department of Geological Sciences, Northwestern University, 1850 Campus Dr., Evanston, IL 60208, USA.

⁵Hawaii Institute of Marine Biology, PO Box 1346, Kaneohe, HI 96744, USA.

¹ Laboratory for Global Marine and Atmospheric Chemistry, School of Environmental Sciences, University of East Anglia, Norwich, Norfolk NR4 7TJ, U.K. (f.hopkins@uea.ac.uk)

² Plymouth Marine Laboratory, Prospect Place, West Hoe, Plymouth, Devon PL1 3DH, U.K.

³ Department of Biological Sciences, University of Essex, Wivenhoe Park, Colchester, Essex CO4 3SQ, U.K.

plankton growth rate vs. temperature (the Eppley curve). By doing so, we provide a conceptual framework for reconciling the results of available experimental manipulations of coccolithophores and a route-map to a simple quasi-empirical relationship for use in carbon cycle models. The implications of an 'Eppley' like calcification formulation for future fossil fuel CO₂ uptake are explored in the GENIE-1 Earth system model.

DAY 4: THURSDAY, 9 OCTOBER

BEYOND NATURAL SCIENCE

A SCIENCE SUMMARY OF THE SYMPOSIUM [I]

Carol Turley, Plymouth Marine Laboratory, U.K.

AN ECONOMIC PERSPECTIVE ON OCEAN ACIDIFICATION AND ATMOSPHERIC CARBON DIOXIDE STABILISATION [I]

Held, Hermann and Ottmar Edenhofer

Ocean acidification represents one of the most significant impacts of a rising CO2 concentration in the atmosphere. Within the upcoming negotiations about a Post-2012 'global climate deal', it should therefore shift the balance towards further CO2 emission mitigation efforts. Here we condense the strain of arguments that are hampering stringent mitigation measures until today and that were academically overcome only recently, summarised in two major reviews: Stern's report to the British government (2006) as well as the latest IPCC report (2007). Both reports outline that 'binding mitigation measures' (such as a 2° or a 450 ppm CO2 target) could be implemented at almost negligible reduction in economic welfare, while at the same time avoiding CO2-induced damages that are potentially an order of magnitude larger. We outline the key shifts in investment streams necessary to build-up emission-free capacities, and sketch structural elements of the 'global deal' that could mobilise the according investment switches. When designing such a deal, it will be necessary to weigh the impacts avoided through mitigation against potential impacts induced by mitigation technologies such as bio-energy, CCS (carbon capture and sequestration) or nuclear energy. We put ocean acidification into that context and argue that the ocean acidification community must increasingly become an active player in the upcoming negotiations on such a 'global deal'. This would facilitate better-informed political decisions as it would stimulate early-on additional research priorities in the ocean acidification community.

Potsdam Institute for Climate Impact Research (PIK), PO Box 60 12 03, 14412 Potsdam, Germany; hermann.held@pik-potsdam.de .

OCEAN ACIDIFICATION- CONNECTING THE SCIENCE TO POLICY [I]

John M Baxter¹ and Dan Laffoley²

Undertaking original scientific research has always been the response to seeking understanding of new issues. By nature scientists are curious animals but once they have identified and answered the questions that interest them, they publish their findings in a peer-reviewed journal and move on to the next question that interests them.

The complexity and range of issues that we face today means that no longer can anyone be expert or even well-informed about everything, and as such there is great danger that important emerging facts

¹ University of Bristol, Bristol BS8 1SS, U.K. (andy@seao2.org),

² Department of Earth Sciences, University of Bristol BS8 1RJ, U.K.,

³ Plymouth Marine Laboratory, Plymouth PL1 3DH, U.K.

about some key issues may be overlooked. Ocean acidification is one such area where we can't afford that to happen. The world is at last more-or-less now aware of the challenges presented by global climate change but to most the consequences of this translates to rising temperatures, melting ice-caps, perhaps increased storminess and changing rainfall patterns, but not ocean acidification which remains one of the largest unrecognised ticking time-bombs.

It is essential that as scientists gain a greater understanding of the mechanisms, rates and implications of this process, this is done in such a way as immediately informs a response by those in the position to act – the decision and policy makers.

We present an approach that has been adopted by EPOCA involving a Reference User Group (RUG) that connects the researchers with a range of relevant people ('the customers') providing the opportunity to interact throughout the lifetime of the project. This helps give the scientists a steer to think about how their research can help answer the burning policy questions, provides an on-going forum to share understanding and to react to changing policy priorities, and provides a platform for better targeted and tailored communications to different audiences who need to know but would never look at the peer-reviewed literature. The UK Marine Climate Change Impacts Partnership (MCCIP) Annual Report Card (ARC) is presented as an example of how complex messages can be distilled down to the headline points and made accessible to policy advisers and government ministers.

¹Scottish Natural Heritage, Silvan House, 3rd Floor East, 231 Corstorphine Road, Edinburgh, Scotland, EH12 7AT, U.K.

²Natural England, Northminster House, Peterborough, England, PE 1 1UA, U.K.

POSTER ABSTRACTS

The abstracts in this section are arranged in alphabetical order by the family name of the first author, whether or not that person is presenting the poster. The presenting author's name is underlined. All posters will be available for viewing throughout the Symposium. There will be poster sessions on Monday and Tuesday evenings. At each of these sessions half of the authors will present their posters. The poster titles are followed by an M (for presentation on Monday evening) or by a T (for presentation in the Tuesday poster session). See the index at the back of this book for a list of all authors.

RECENT CHANGES IN THE INORGANIC CARBON SYSTEM IN A LARGE, COMPLEX ESTUARY (PUGET SOUND, WASHINGTON, USA) M1

Alin, Simone R.¹, Richard A. Feely¹, Christopher L. Sabine¹, Jan Newton², Skip L. Albertson³, and Cynthia G. Peacock¹

Puget Sound is a large estuary complex with a diversity of habitat types and 10,000 rivers and streams draining into it. Circulation within the basin and between the basin and the open ocean is limited by relatively shallow sills, making Puget Sound quite sensitive to changes in freshwater and nutrient inputs related to changes in land use, streamflows, climate, and coastal ocean conditions. Rising regional air temperatures, decreasing freshwater inputs, and human alterations to the coastline during the 20th century have likely changed the carbon distributions in the Sound, but historical carbon data are very limited. In 2008, we conducted the first extensive high-quality surveys of inorganic carbon distributions in the Puget Sound. To understand how the carbon system has changed over time, we have analyzed the Puget Sound Assessment and Monitoring database (1989-2008) for changes in water chemistry that may have implications for the Sound's inorganic carbon system and food web. Over the period of record, average annual deep water temperatures have increased by 2°C, and salinity has decreased by 2. A preliminary model suggests that the effects of these changes may correspond to a reduction in aragonite and calcite saturation levels in the deep water environment. Using discrete samples collected for dissolved inorganic carbon, alkalinity, and oxygen during the 2008 survey cruises, we are calibrating this model to more accurately represent changes in the inorganic carbon system associated with climate change.

PHYSIOLOGY DRIVES CORAL-REEF COMMUNITY SHIFTS UNDER CLIMATE CHANGE T1

Anthony, Ken and Guillermo Diaz-Pulido

The fate of coral reefs under climate change is of global concern. Our knowledge of how coral-reef organisms respond to thermal stress and ocean acidification is growing, but we know little about the linkages between organism-level stress responses and the behaviour of reef communities. The dynamics of benthic reef communities and their implications for reef resilience are mostly studied from the perspective of top-down control processes (e.g. algal grazing) and the frequency and severity of major physical disturbances (e.g. cyclones). Here, we analyse the relative importance of physiological susceptibility to high-CO₂ scenarios in driving benthic community dynamics. We use the functional responses of growth and mortality risk of corals and macroalgae to a combined temperature-acidification gradient as input into a community dynamics model and analyse changes in equilibrium compositions. Relative response strengths of coral and algal growth and mortality were

¹Pacific Marine Environmental Laboratory, National Oceanic and Atmospheric Administration, Seattle, Washington 98115, USA (simone.r.alin@noaa.gov).

²Applied Physics Laboratory, University of Washington, Seatlle, Washington 98195, USA.

³Washington Department of Ecology, Olympia, Washington 98504, USA.

derived from experimental studies and partly from varying stress responses in set of sensitivity analyses. Model projections demonstrated that the location of thresholds for shifts between coral and macroalgal dominated states depends strongly on how sensitive coral and algal growth rates and mortality risks are to climate change. Interestingly, climate-change effects interacted synergistically with grazing intensity and nutrient loading in driving the system to algal dominance. High sensitivity of some macroalgal species to acidification can shift reef systems to barrens as an alternate stable state. Our results suggest that bottom-up processes from acidification and thermal stress are a stronger determinant of future reef compositions than top-down control processes.

Centre for Marine Studies, ARC Centre of Excellence for Coral Reef Studies, The University of Queensland, St. Lucia, Queensland 4072, Australia

EFFECTS OF CO₂ SEEPAGE ON SOLUBILITY AND TRANSPORT OF TRACE ELEMENTS IN SEAWATER AND SEDIMENT M2

Ardelan, Murat V. and Eiliv Steinnes

Two lab-scale chambers have been developed to study the effects of CO_2 leakage on trace metal/elements biogeochemistry in seawater and the sediment-water interface

During the initial phase of CO_2 seepage (15 days) "dissolved" (< 0.2 µm) concentrations of all trace transition metals (Al, Cr, Mn, Fe, Co, Ni, Cu, Zn, Cd and Pb) increased significantly (> 50 %) in the water. During a second phase of additional 10 days with the same sediment but replenished seawater Al, Cr, Cd, and Zn were partly removed from the water column in the CO_2 chamber, Cu still increased but at reduced rates, whereas Pb increased faster than during the first period. Dissolved fractions of all other 20 elements increased substantially during early CO_2 seepage except Sc, Ga, Ge, As, Sn and Cs. DGT-labile fractions (Easily diffusive forms through the DGT sampling) of all trace transition metals increased during early CO_2 seepage, and continued to increase during the second phase except for Cu and Zn. In the sediment pore water all trace metals, except Mn, increased drastically during CO_2 seepage. DGT labile Sc, Ti, V, As, Y, Sn, Cs, La, Ce, Pr, Sm, Tb, Th and U increased with CO_2 seepage in both sediment pore water and in the overlying water.

There are distinct patterns among different trace elements regarding effects of CO_2 seepage. More "realistic" experiments with the high-pressurize TiO_2 tank soon ready for use will definitely yield give important information on CO_2 effects on bioactive and toxic trace element distributions in the sediment – seawater system.

Norwegian University of Science and Technology, Department of Chemistry, NO 7491, Trondheim, Norway. (murato@nt.ntnu.no)

DMS AND DMSP CYCLING IN A CHANGING MEDITERRANEAN SEA- AN ACIDIFICATION APPROACH T2

<u>Avgoustidi, V.¹</u>, N. Michalopoulos², U. Christaki³, E. Krasakopoulou¹, S. Economou¹, A. Konstantinopoulou¹, and E. Papathanassiou¹

Changing seawater pH is likely to have a major impact on the marine sources of climatically active gases such as dimethyl sulphide (DMS), as it has been shown previously. The production of DMS by phytoplankton and its atmospheric oxidation products have been coupled to climate forcing. This work presents a first attempt to investigate the link between increasing pCO₂ and DMS concentrations as well as bacterial protein production and abundance, in the Eastern Mediterranean Sea; an area characterised by extreme oligotrophy and phosphorus limitation. The Eastern Basin of the Mediterranean Sea is highly sensitive to climate change and anthropogenic activity and is considered as one of the most oligotrophic areas in the world, where heterotrophic bacterial production is tightly coupled to primary productivity. The current study describes the potential effect that increasing atmospheric CO₂ concentrations may have on the DMS concentrations and bacterial biomass and activity, derived from natural phytoplankton communities. Acidification experiments are presented, in

which natural phytoplankton assemblages have developed in enriched CO₂ environments and current CO₂ concentrations. Recent pCO₂ measurements carried out in the area where water has been collected from indicate that throughout the year, surface seawater is undersaturated with respect to atmospheric pCO₂. We report changes in the concentrations of DMS and its cellular precursor DMSP, in response to this CO₂ enrichment and investigate the role of bacteria on the DMS and DMSP cycling in a high CO₂ world.

SATURATION STATE OF SEA WATER FOR CALCITE AND ARAGONITE IN CANADIAN ARCTIC ARCHEPELAGO AND THE LABRADOR SEA M3

Azetsu-Scott, Kumiko¹, David Slauenwhite¹ and Philip Yeats²

Channels of the Canadian Arctic Archipelago (CAA) and the Hudson Bay region provide the main pathway for the flow from the Arctic Ocean to the North Atlantic. Rapidly changing physical conditions in the Arctic, such as hydrological cycles, sea water temperature and reduction of ice cover impact biogeochemical cycles in the Arctic and propagate through the CAA and the Hudson Bay to the Labrador Sea, where deep convection occurs in winter and influences the global thermohaline circulation. Saturation state of seawater with respect to calcite and aragonite was calculated from dissolved inorganic carbon and total alkalinity collected from the CAA to the Labrador Sea in 2003-2004. Transects include Nares Strait, Barrow Strait, Baffin Bay, Hudson Bay, Hudson Strait, Davis Strait, and the Labrador Sea. While Barrow Strait was supersaturated for both calcite and aragonite through the entire water column (deepest 280m), shallow saturation depths, at 150m for calcite and at 75m for aragonite, were observed in the Hudson Bay. Saturation depths for aragonite in Davis Strait and in the Labrador Sea were 800m and 2500m, respectively. Variation in saturation depths in different transects is described from water mass structure, circulation and organic matter production/respiration.

SHORT-TERM RESPONSE OF THE COCCOLITHOPHORE *EMILIANIA HUXLEYI* TO AN ABRUPT CHANGE IN SEAWATER pCO_2 T3

Barcelos e Ramos, J., M. N. Müller and U. Riebesell

The response of the coccolithophore *Emiliania huxleyi* to future CO₂ increase is well documented in acclimated cultures where cells are exposed to the CO₂ conditions for several generations before the experiment. However, the time necessary to reach a new physiological "equilibrium" (acclimation) remains unknown. Here we show that *Emiliania huxleyi's* short-term response (hours to 1 day) to increasing CO₂ is similar to previous studies with acclimated cultures under comparable conditions. Indeed at CO₂ levels from glacial (190 µatm) to year 2100 (750 µatm), calcification decreased and organic carbon fixation increased already 8 h after exposing the cultures to the changed CO₂ conditions. This led to a decrease in the ratio of CaCO₃ to organic carbon production. Our results show that with the change in CO₂ concentration a new physiological "state" is established in a matter of hours, apparently independent of cellular division. If these results apply to other phytoplankton species, it has implications for the interpretation of mesocosm studies and on board incubations, where often it is not feasible to allow for a pre-conditioning phase before starting experimental incubations.

¹Hellenic Centre for Marine Research, Anavissos, Greece (vavgoustidi@ath.hcmr.gr)

²University of Crete, Heraklion, Crete, Greece.

³Universite du Littoral, Cote d'Opale, France

¹ Ocean Sciences Division, Department of Fisheries and Oceans, Bedford institute of Oceanography, Dartmouth, Nova Scotia, B2Y 4A2, Canada (Azetsu-ScottK@mar.dfo-mpo.gc.ca)

² Ecosystem Research Division, Department of Fisheries and Oceans, Bedford institute of Oceanography, Dartmouth, Nova Scotia, B2Y 4A2, Canada.

Leibniz Institute of Marine Sciences, IFM-GEOMAR Düsternbrooker Weg 20, 24105 Kiel, Germany. jramos@ifm-geomar.de

OCEAN ACIDIFICATION AND THE DEMISE OF BRACHIOPODS M4

Barry, James P., Eric F. Pane, Kurt R. Buck, Chris Lovera, and Christina Tanner

Brachiopods, commonly called lamp-shells, were a dominant element of Paleozoic seas when perhaps 30,000 species thrived. During the Permian-Triassic extinction brachiopod diversity dropped to ~300 species, perhaps related to ocean acidification, hypoxia, and other stresses during this event. The survivors diversified slightly during the Mesozoic, but once again were overwhelmed during the mass extinction at the end of the Cretaceous. Today, brachiopods are a somewhat rare member of marine ecosystems, found principally in intertidal ~1 km depths throughout the world ocean.

Although brachiopods are thought to be susceptible to stresses caused by ocean acidification, including reduced calcification, respiratory stress, acidosis, and metabolic depression, no studies have investigated their tolerance to ocean acidification. We examined the physiology, survival, and growth of brachiopods (*Laqueus* sp.) common on the continental shelf and upper continental slope in the eastern Pacific, in response to various levels of environmental hypercapnia (ocean acidification). Under mild (~0.2 pH units) to severe (~1.0 pH units) environmental hypercapnia, the internal fluids of brachiopods were mostly isotonic with seawater, indicating that they have little capacity to regulate internal fluid chemistry, or buffer acidosis caused by ocean acidification. Metabolic rates (oxygen consumption) were initially elevated under high pCO2 / low pH treatments, compared to controls, but slowed with continued exposure. Longer-term studies are evaluating the effects of hypercapnic stress on growth and survival. Our results confirm that brachiopods suffer reduced metabolic function under stresses associated with ocean acidification, which likely lead to death and perhaps species extinctions.

Monterey Bay Aquarium Research Institute, 7700 Sandholdt Road, Moss Landing, CA 95039, USA (barry@mbari.org)

VULNERABILITY OF SOUTHERN OCEAN PTEROPODS TO ANTROPOGENIC ACIDIFICATION T4

Bednarsek, Nina^{1,2}, Geraint Tarling ^{1,} Dorothee Bakker², and Sophie Fielding¹

Uptake of anthropogenic carbon dioxide (CO₂) is changing the carbonate chemistry in the Southern Ocean. It has been suggested that marine calcifiers, among them pteropods, will be affected by the resulting decrease in ocean pH (Kleypas et al., 2005). Orr et al. (2005) demonstrated rapid shell dissolution in low pH waters, however a comprehensive study of the effect of ocean acidification on Southern Ocean pteropods is lacking. Well advanced research is ongoing to determine the life cycle of Limacina helicina by studying net samples from historic James Clark Ross cruises (early 90's -2008). Additional net samples from the JR177 (2008) cruise in the Scotia Sea will be combined with vertical profiles of carbonate parameters, thus allowing the study of pteropod abundance versus the aragonite saturation state. Furthermore, high CO₂ experiments were carried out on cruise JR177, which focused on the susceptibility of the pteropod larval stages to ocean acidification. Sampled individuals of Limacina helicina were incubated at different levels of the partial pressure of CO₂ (375, 500, 750 and 1200 p.p.m.v.) for up to 13 days to mimic ocean acidification. Upon the return of the samples to the U.K., the organisms will be analysed for the effect of acidification on their morphology and mineral composition of the shell. Water samples were also preserved for analysis of total alkalinity (TA) and total dissolved inorganic carbon (TCO₂). From these studies the capability of pteropod larvae to withstand changes in carbonate chemistry will be assessed.

¹British Antarctic Survey, Madingley Road, High Cross, Cambridge CB30ET, U.K. (Bednarsek, N: nindna@bas.ac.uk, Tarling, G: gant@bas.ac.uk, Fielding, S: sof@bas.ac.uk)

²University of East Anglia, School of Environmental Sciences, Norwich NR4 7TJ, U.K. (Bakker, D: D.Bakker@uea.ac.uk)

THE CHALLENGES OF MODELLING ECOSYSTEM RESPONSE TO OCEAN ACIDIFICATION. M5

Blackford, Jerry

The chemistry of ocean acidification is relatively predictable but the precise response of the many sensitive ecological and biogeochemical processes remains speculative. Experiments have revealed diverse responses for different species although few species have been studied. Over the same timescale as acidification, global warming will alter the physical drivers of the marine system, prediction therefore requires an integrated multi-driver approach. Modelling provides our only predictive tool but is limited by contentious parameterisations, a lack of evaluation data and complexity. Given these uncertainties, it is important not to be overly deterministic with models, however models do have an important role. These include: predicting carbonate system parameters and carbonate saturation states under different emission scenarios, assessing the sensitivity to parameter uncertainty and model structure, producing probabilistic ensemble scenarios rather than deterministic assessments, testing if a given response is by itself capable of altering key system properties, assessing if an acidification response might be affected by changing climate or other drivers, asking how do process responses identified in (mono-)culture type studies express within a virtual complex ecosystem and finally as provocateurs of debate and experimentation. To illustrate these approaches, results from 3D simulations of the North West European Shelf will show predictions for the carbonate system, an assessment of the vulnerability of the nitrogen cycle and benthic coupling due to the inhibition of nitrification, an illustration of how phytoplankton species specific carbon uptake rates might provoke changing community structure and an examination of the sensitivity of acidification response to changing climate.

Plymouth Marine Laboratory, Prospect Place, Plymouth, PL1 3DH, U.K. (jcb@pml.ac.uk).

KIEL CO₂ MANIPULATION EXPERIMENTAL FACILITY (KICO2) T5

Bleich, Markus¹, Frank Melzner², Claas Hiebenthal², Helgi Mempel², Kai Schulz², Ulf Riebesell², Martin Wahl², Frank Sommer², Ulrich Sommer², Armin Form², Uwe Piatkowski², Reinhold Hanel², Dieter Piepenburg³, Michael Spindler³, Anton Eisenhauer², Andrea Franke², Volker Möller⁵, Gunnar A. Baumert⁴, and Catriona Clemmesen²

We will present the technical details and our first experiences with a new CO₂-manipulation facility, set up at IFM-GEOMAR in March 2008. The system adds pure CO₂ to pressurized air to create 5 different CO₂-air mixtures (380-4,000 ppm of CO₂), which are distributed into 6 different temperature constant rooms at gas flow rates of up to 1000 l/h per room and per CO2 concentration. Precise addition of CO₂ is achieved via computer controlled valve systems (red-y smart series) that respond to variations in gas flow according to the gas demand by the experiments. CO₂ flow is detected by pressure and temperature independent thermal measurement of mass. High precision of final CO₂ values is achieved within 1-100% of maximal gas flow rate. Actual values and set points are continuously monitored and documented online via PC interface. The facility is completed by a security and alarm system. The gas mixtures are used to equilibrate different types of experimental setups: (1) large flow-through Baltic Sea systems for benthic invertebrate long-term incubations, (2) small flow-through systems for short- and long-term incubations of Baltic- and North Sea invertebrates, (3) small-scale larval incubation systems for simultaneous manipulation of CO₂ and temperature. All types of experimental setups will be described in detail and quantitative information of crucial parameters (e.g. water and gas flow rates, TA, DIC, pH, experimental animal biomasses, and nitrogenous waste accumulation) will be given and discussed. This contribution is an effort towards establishing guidelines for highly controlled and reproducible CO₂ manipulation experiments.

OCEAN ACIDIFICATION AND HARMFUL ALGAL BLOOMS M6

Boisson, Florence, Jean-Louis Teyssié, François Oberhansli, and Ross Jeffree

Scientific knowledge on biomineralization suggests that changes in pH associated with ocean acidification will have consequences on the physiology of planktonic calcifiers such as coccolithophores, foraminifera, and pteropods, modifying the equilibrium within plankton communities. Then, in the context of ocean acidification, we can ask what potential is there for an increase in non-calcifying plankton that is responsible for Harmful Algal Blooms (HABs), which strongly affect seafood resources worldwide? Harmful algae physiology and toxin content may be impacted by changes in seawater chemistry, as well as simultaneous warming and associated variations in metal bioavailability and nutrient supply. Here we will present plans to use experimental radiotracer-assay methods developed at the Radioecology Laboratory of the IAEA-Marine Environment Laboratories to contribute to future studies of harmful algal cultures (namely those responsible for Paralytic Shellfish Poisoning) that need to be designed to test the hypothesis that HABs will increase in a future High-CO2 world.

Marine Environment Laboratories, International Atomic Energy Agency, 4 Quai Antoine Ier, MC98000 Monaco, Principality of Monaco (F.Boisson@iaea.org)

INCREASED CONTENT OF ${\rm CO_2}$ IN THE ATMOSPHERE CAN REDUCE PRIMARY PRODOCTION IN THE OCEAN T6

Børsheim, Knut Yngve

Recently it has been showed that the average C:N ratio may increase in a high CO₂ world, which may imply an increased availability of organic carbon in the future ocean. This may stimulate heterotrophy, but heterotrophic prokaryotes are not only dependent on DOC for their life support, they also compete with the primary producers for mineral nutrients. In our group based in Bergen, Norway, we have conducted several studies on the flow of carbon in pelagic plankton, using both laboratory culture experiments and field work including mesocosm perturbation methodology. In a series of experiments both mineral nutrients and glucose were added at a suite of combinations. Glucose was envisaged as a proxy for labile dissolved organic carbon. The results showed that when the system was nutrient limited, adding more organic carbon resulted in smaller total accumulation of organic carbon in the system. We argue that stimulation of the heterotrophs under mineral nutrient limitation represents a cost to primary producers, and under such conditions increased availability of organic carbon decreases the total production in the system.

Institute of Marine Research, P.b. 1870 Nordnes, NO-5817 Bergen, Norway (yngve.borsheim@imr.no)

¹ Institute of Physiology, CAU Kiel, Olshausenstraße 40, 24098 Kiel, Germany (m.bleich@physiologie.uni-kiel.de).

² IFM-GEOMAR, Leibniz Institute of Marine Sciences, Wischhofstraße 1-3, 24148 Kiel, Germany (cclemmesen@ifm-geomar.de)

³ Institute for Polar Ecology, CAU Kiel, Wischhofstraße 1-3, 24148 Kiel, Germany

⁴ HTK Hamburg GmbH, Woelckenstraße 11, 22393 Hamburg-Sasel, Germany (gab@htk-hamburg.com)

⁵ Linde AG Geschäftsbereich Linde Gas, Fangdieckstraße 75, 22547 Hamburg Germany (Volker.Moeller@de.linde-gas.com)

HIGH FREQUENCY MONITORING OF THE PARTIAL PRESSURE OF CO_2 (pCO₂) USING A CARIOCA SENSOR ON A MAREL BUOY IN A TEMPERATE COASTAL ECOSYSTEM (2003-2008) – A TOOL FOR INVESTIGATING THE OCEAN IN A HIGH CO_2 WORLD M7

<u>Bozec, Yann¹</u>, Liliane Merlivat², Laurence Beaumont³, Théodore Danguy³, Antoine Guillot³, Michel Répécaud⁴, Emilie Grossteffan⁵, Eva Bucciarelli⁶, Jacques Guillou⁶, Stéphane Blain⁷, and Paul Tréguer⁶

Long-term monitoring of the marine carbon chemical species is necessary to assess the chemical and biological modifications occurring in the ocean in a "High CO₂ World". Coastal marine ecosystems are directly impacted by human activities and are crossing a threshold of changing from their preindustrial state, during which ocean margins are widely viewed as heterotrophic and a CO₂ source, to a current or future state as a CO₂ sink.

The CARbon Interface Ocean Atmosphere (CARIOCA) sensor allows for both long term and high frequency measurements of the partial pressure of CO₂ (pCO₂). The CARIOCA sensor is therefore an excellent tool for investigating the high variability and the evolution of pCO₂ in coastal environments. Here we present high-frequency pCO₂ data recorded for 5 years during the first deployment of a CARIOCA sensor on a MAREL buoy in the surface waters of a temperate coastal ecosystem, the Bay of Brest, which is impacted by both coastal and oceanic variability. High frequency measurements allowed for the quantification of the diurnal, tidal and seasonal variability in the assessment of the annual CO₂ air-sea fluxes. The preliminary results indicate that biological activity is the main process controlling the pCO₂ variability in surface waters on a seasonal time-scale. On a shorter scale, the tidal and diurnal cycle are shown to be responsible for high pCO₂ variability. The 5 years of investigation revealed that the surface waters of the Bay were near equilibrium with the atmosphere and that the inter-annual variability was small.

SYNERGISTIC EFFECTS OF CO₂ CONCENTRATION AND TEMPERATURE ON OXYGEN EXCHANGE IN CORALS T7

Buxton, Lucy¹, M. Badger² and P. Ralph¹

In the last 100 years, human activities have contributed considerably to atmospheric CO₂ emissions. CO₂ is recognized as a major greenhouse gas and causal links have now been drawn between atmospheric CO₂ and increases to global mean temperatures. However, high atmospheric CO₂ levels have recently been shown to cause chemical changes in oceanic surface waters, including increased concentrations of dissolved CO₂, decreased bicarbonate ion concentration (CO₃²). Impacts of altered carbon concentration are in addition to that of temperature alone, and the negative effects on oceanic flora and fauna will be substantial. Amongst the systems likely to be impacted by these biochemical changes, coral reef ecosystems are thought to be especially vulnerable because of their complex carbon budgets linked to photosynthesis and calcification.

The interactions of temperature and C_i concentration remain poorly understood because physiological responses have been primarily examined by manipulating one parameter at a time. This has been exacerbated by the fact that previous investigations have relied on either oxygen evolution rates, or measurement of effective quantum yield of PSII as an indicator of photosynthetic functionality. Used for the first time on corals, membrane inlet mass spectrometry (MIMS) allows the continuous measurement of gas exchange (O_2 evol and O_2 uptk) and chlorophyll fluorescence to occur non-invasively Results are presented using combined measurement photosynthesis determined by of O_2

¹UMR 7144 CNRS-UPMC-INSU, Station Biologique de Roscoff (bozec@sb-roscoff.fr)

²LOCEAN, Université Pierre et Marie Curie, Paris

³Division Technique INSU-CNRS

⁴IFREMER, Centre de Brest

⁵UMS 3113 CNRS-UBO, Institut Universitaire Européen de la Mer (IUEM), Brest

⁶UMR 6539 CNRS-UBO, IUEM, Brest

⁷UMR 7621 CNRS-UPMC, Observatoire Océanologique de Banyuls sur Mer

evolution, O_2 uptake and chlorophyll *a* fluorometry of the branching coral *Pocillopora damicornis* exposed to a range of C_i concentrations (0.1 – 12 mM) at 26 and 30 °C over short-term incubations.

¹Department of Environmental Sciences, Institute for Water and Environmental Resource Management, University of Technology Sydney, Sydney, NSW 2007, Australia (lucy.buxton@uts.edu.au)

²ARC Centre of Excellence in Plant Energy Biology, Molecular Plant Physiology Group, Research School of Biological Sciences, The Australian National University, Canberra, ACT, Australia

IMPACT OF SEAWATER ACIDIFICATION ON SEA URCHIN SKELETAL GROWTH M8 Catarino, Ana I. and Philippe Dubois

Sea urchins are particularly exposed to seawater acidification due to the nature of their well developed high-magnesium calcite skeleton. This is formed trough an amorphous calcium carbonate precursor form whose solubility far exceeds that of aragonite. As sea urchins are keystone species in numerous ecosystems, the adverse effects they might experience will have severe ecological repercussions. In this study we assessed the impact of seawater acidification on skeletal growth in both larvae and adults of the temperate sea urchin *Paracentrotus lividus*.

Experiments were carried out at controlled temperature and lower pHs were obtained by bubbling CO_2 in seawater. The total alkalinity and pH (seawater scale) were measured and the pCO_2 and total inorganic carbon calculated, as well as the magnesium-calcite saturation state. The larvae were grown in pHs between 8.0-6.8 until the echinopluteus stage and their development was characterized using embryotoxicity protocols to distinguish normal and altered morphologies. The spicules development was quantified. At lower pHs significantly more larvae showed an abnormal morphology and a reduced size, going together with frailer spicules. Adult sea urchins were grown at pHs between 7.8-7.2. Spine regeneration was experimentally induced and regenerate sizes were measured using microscopy techniques. Their magnesium content was analysed and alteration of mechanical properties was evaluated.

Available data already indicate that both stages of the sea urchin life cycle are affected by sea water acidification. This suggests that ecosystems structured by sea urchins will be seriously affected by ongoing ocean acidification.

Laboratoire de Biologie Marine, Université Libre de Bruxelles, CP 160/15, avenue F.D. Roosevelt 50, 1050 Bruxelles, BELGIUM (ana.catarino@ulb.ac.be)

EXPLORING THE RESPONSE OF *IN SITU* COCCOLITHOPHORE POPULATIONS IN THE NORTH EAST ATLANTIC TO ACIDITY AND CALCITE SATURATION GRADIENTS. T8

<u>Charalampopoulou, Anastasia</u>¹, Cynthia Dumousseaud¹, Alex Poulton¹, Mike Lucas², Darryl Green¹, Toby Tyrrell¹ and Eric Achterberg¹

Ocean acidification is expected to have a negative impact on marine calcifiers, specifically on coccolithophores. Coccolithophores are of particular importance as they comprise a great proportion of pelagic biogenic calcification (50-80%). Moreover, calcite is considered to be an important ballast material for organic carbon transport from the surface to the deep ocean and therefore plays an important role in the global carbon cycle.

The fate of coccolithophores in a future acidic ocean has been intensively researched during the last few years. Most of our knowledge comes from laboratory experiments on monospecific cultures, the results of which are sometimes contradictory, and research on natural populations has been limited. To investigate how natural assemblages might respond to ocean acidification we simultaneously sampled natural coccolithophore populations and the ocean carbonate system during a cruise to the eastern Iceland basin in July-August 2007. Specifically, we made simultaneous *in situ* measurements of particulate inorganic carbon (PIC), particulate organic carbon (POC) and calcification rates (ρ PIC),

and we used Scanning Electron Microscopy (SEM) to assess coccolithophore abundance and morphology. We also measured dissolved inorganic carbon (DIC) and alkalinity (TA) to derive other parameters of the carbonate system and specifically the calcite saturation state of the study area. This work describes the observed relationships between the different biological parameters above, one of them being a positive linear relationship between PIC and ρ PIC, and also between them and the carbonate system chemistry.

¹National Oceanography Centre, Southampton, European Way, Southampton, U.K., SO14 3ZH (a.charalampopoulou@soton.ac.uk)

DISPERSION OF CO₂ LEAKED FROM SEABED IN SEAWATER M9

Chen, Baixin¹, Peter Brewer², and Masahiro Nishio³

We have developed a numerical model of the dispersion of CO₂ droplets/bubbles in seawater to simulate the fate of CO₂ leaked or released from seabed naturally or artificially.

The model has been applied to predicting the dynamics of leaked CO_2 in seawater at various depths from 10-800m (temperature from 5 °C to 25 °C) and for initial droplet/bubble sizes from 3.0 to 40.0mm diameter. A diagram of CO_2 ascending distance vs dissolution time is obtained from model simulations. It is found that CO_2 droplets ascend at a mean speed of 11cm/sec and a mean shrinking rate of 7.0 x10⁻³ mm/sec in diameter approximately if leaked from a deep-ocean (800–1000m) source. This speed and shrinking rate increase to 16 cm/sec and 30 x10⁻³ mm/sec at middle deep ocean (500–650m) and finally reach to 22 cm/sec and 0.2 mm/sec at shallow ocean (<150m). At shallow depths, a CO_2 bubble with initial diameter of 40 mm released from 40 m depth might be just dissolved completely before reaching the sea surface, while a bubble with size smaller than 40mm leaked from seabed at the same depth will be fully dissolved.

The two-phase turbulent plume model predicts that the pH change in seawater due to CO_2 dissolution is directly proportional to the leakage rate and inversely proportional to initial droplet/bubble size (do) and ocean current speed (Vc). At a leakage rate of 1.0 kg/sec, do=8.0mm, and Vc =2.5cm/sec, the maximum pH change is -1.6. This pH change increases to -2.5 when leakage rate increases to 10kg/sec and reduces to -1.2 when leakage rate decreases to 0.1kg/sec, respectively. The mean CO_2 enriched seawater volume developed from CO_2 leakage position in a turbulent ocean is also simulated. The results show that the volume with pH changes larger than -0.1 could increase up to 2.0×10^6 m³ within 30 minutes of onset of the source.

² Monterey Bay Aquarium Research Institute, Moss Landing, CA 95039, USA (brpe@ mbari.org)

BIOGEOCHEMICAL INVESTIGATIONS OF COCCOLITHOPHORE BLOOMS ALONG THE CONTENTAL MARGIN OF THE NORTHERN BAY OF BISCAY: HIGHLIGHTS OF THE PEACE PROJECT T9

<u>Chou, Lei</u>¹, Jérôme Harlay¹, Caroline De Bodt¹, Nathalie Roevros¹, Alberto V. Borges², Kim Suykens², Bruno Delille², Koen Sabbe³, Nicolas Van Oostende³, Anja Engel⁴, Judith Piontek⁴, Corinna Borchard⁴, Nicole Händel⁴, Sabine Schmidt⁵, and Steve Groom⁶

Recent studies have demonstrated that changing ocean chemistry due to ocean acidification poses a growing threat for marine organisms such as corals, coccolithophores and many others that form calcareous skeletons. Its biogeochemical feedbacks and impact on the oceanic carbon cycle are yet to be quantified. Coccolithophores are the major calcifying phytoplankton in the sub-polar and temperate regions of the world's ocean. They produce furthermore Transparent Exopolymer Particles (TEP), which are known to promote aggregate formation. Combined with the CaCO₃ ballast effect, large-

²Department of Zoology, University of Cape Town, Cape Town, South Africa.

¹ Heriot-Watt University, Edinburgh, EH14 4AS, U.K. (b.chen@hw.ac.uk)

³ National Institute of advanced Industrial Science & Technology, Tsukuba, 305 8564, Japan (m.nishio@aist.go.jp)

scale coccolithophore blooms could thus contribute to the export of organic carbon to deep waters on relatively short time scales. During the PEACE project, funded by the Belgian Federal Science Policy Office, we have conducted in three consecutive years interdisciplinary biogeochemical surveys, assisted by remote sensing, along the continental margin of the Northern Bay of Biscay where coccolithophore blooms dominated by *Emiliania huxleyi* are frequently and recurrently observed. Rates of various processes governing the coccolithophore ecosystem dynamics have been determined. They include for example primary production, calcification, pelagic and benthic respiration and dissolved esterase enzyme activity. Particle dynamics and organic carbon export have also been assessed, based on thorium isotope measurements. The data are complemented by other relevant biogeochemical parameters such as chlorophyll *a*, HPLC-pigments, TEP, nutrients, and particulate organic and inorganic carbon. The key results will be presented and discussed in attempt to elucidate the oceanic coccolithophore ecosystem response in a high-CO₂ world.

¹Laboratoire d'Océanographie Chimique et Géochimie des Eaux, Faculté des Sciences, Université Libre de Bruxelles, B-1050 Brussels, Belgium (Lei.Chou@ulb.ac.be)

THE EFFECT OF pCO₂ ON THE EGG DEVELOPMENT AND THE CONDITION OF NEWLY-HATCHED LARVAE OF ATLANTIC HERRING (CLUPEA HARENGUS L.) M10 Clemmesen, Catriona and Andrea Franke

The study addresses the effect of hypercapnia (elevated pCO2) on the early development of the Atlantic herring (Clupea harengus L.), the most sensitive stages in the life history of a fish. Eggs of the Atlantic herring were fertilized and incubated in artificially acidified seawater (pH 7.67, 7.49, 7.33, 7.28 and 7.05) and a control treatment (pH 8.08) until the main hatch of herring larvae occurred. The development of the embryos was monitored daily and newly-hatched larvae were sampled to analyze their morphometrics and their condition by measuring the RNA/DNA ratios. Elevated pCO2 neither affected the fertilization rate nor the embryogenesis, the mortality rate or hatch rate. Furthermore the results showed no linear relationship between pCO2 and yolk sac area, dry weight and total length of the newly-hatched larvae. For pCO2 and RNA/DNA ratio, however, a significant positive linear relationship was found. The RNA concentration at hatching time was lowered, which consequently should lead to a reduced protein synthesis rate. The results indicate that an increased pCO2 can affect the metabolism of fish embryos negatively and therefore further somatic growth of the larvae could be reduced. An additional experiment was conducted using controlled air CO2 mixture with a newly developed automated CO2 manipulation system with concentrations of 380, 560, 870, 1120 and 1400ppm. The effects on the condition of herring larvae at hatch will be analysed and compared.

Leibniz Institute of Marine Sciences (IFM-GEOMAR), Düsternbrooker Weg 20, 24105 Kiel, Germany, cclemmesen@ifm-geomar.de

RESPONSE OF MEDITERRANEAN AND ARCTIC PTEROPODS TO OCEAN ACIDIFICATION T10

Comeau, Steeve¹, Gaby Gorsky¹, Jean-Louis Teyssié², Ross Jeffree², and Jean-Pierre Gattuso¹

Thecosome pteropods (molluses) can play an important role in the food web of various ecosystems. For example, polar species are reported to be an important food source for herring, salmon and whales. Ocean acidification driven by the increase of pCO2 could have an important impact on these

²Unité d'Océanographie Chimique, Université de Liège, B-4000 Liège, Belgium

³Protistologie & Aquatische Ecologie, Universiteit Gent, B-9000 Gent, Belgium

⁴Alfred Wegener Institute for Polar and Marine Research, D-27515 Bremerhaven, Germany

⁵Environnements et Paléoenvironnements Océaniques (UMR 5805 EPOC – OASU), Université Bordeaux 1, F-33405 Talence, France

⁶Remote Sensing Group, Plymouth Marine Laboratory, Plymouth PL1 3DH, United Kingdom

animals because they harbour an aragonitic shell which was shown to be very sensitive to changes in the carbonate chemistry. Experiments are carried out in the NW Mediterranean sea (*Cavolinia inflexa*, *Limacina inflata*) and in Spitzbergen (*Limacina helicina*). Specific attention is paid to juveniles which could be the most sensitive to a decrease in pH. Despite the fact that pteropods are notoriously difficult to maintain in the laboratory, specimens have been maintained successfully several weeks under controlled conditions of pCO2 (380 and 760 ppm) and temperature, and precipitation of CaCO3 was observed. New culture conditions are being tested to keep them alive in the laboratory for longer periods of time. The fluorochrome calcein and 45Ca are useful tools to investigate calcification. The latest results of the on-going perturbation experiments will be presented.

¹Laboratoire d'Océanographie de Villefranche, CNRS-Université Pierre et Marie Curie-Paris 6, 06234 Villefranche-sur-Mer Cedex, France. comeau@obs-vlfr.fr

SUBANTARCTIC WATER – NO EVIDENCE OF A CO₂ INCREASE. M11 Currie, Kim¹, Keith Hunter² and Malcolm Reid²

The Southern Ocean is known to be a sink for atmospheric CO_2 , however models do not reproduce the observations well, the processes determining the carbonate chemistry in this area are not yet well understood. Inverse models based on atmospheric CO_2 data estimate that the Southern Ocean sink is decreasing due to greater mixing of high carbonate deeper waters to the surface.

pCO₂ and pH in subantarctic water have been directly measured bi-monthly for the last ten years as part of a time series transect programme off the east coast of the South Island of New Zealand (171.50 $^{\circ}$ E 45.85 $^{\circ}$ S). The pCO₂ seasonal cycle has a mean value of 360 \Box atm, with an amplitude of 10 \Box atm, the maximum occurring in early spring. This is out of phase with the effect of changing seawater temperature, biological activity is the main determinant of the seasonal cycle in these waters. There has been no apparent long-term change in either the pCO₂ or the pH since 1998. This has resulted in an increase in the air-sea pCO₂ difference due to the increasing atmospheric carbon dioxide concentration, and thus there has been an increase in the magnitude of the sink for atmospheric carbon dioxide in these waters.

INFLUENCE OF FUTURE CO₂ CONCENTRATIONS ON GROWTH AND NITROGEN FIXATION OF THE BLOOM FORMING CYANOBACTERIUM NODULARIA SPUMIGENA T11

Czerny, Jan T. B., Joana Barcelos e Ramos, and Ulf Riebesell

Anthropogenic CO₂ emissions are resulting in elevated CO₂ concentrations and acidification in the surface ocean. Phytoplankton physiological responses to these chemical alterations may influence ocean primary production and thereby impact pelagic ecosystems and biogeochemical cycling. Diazotrophic cyanobacteria are able to fix carbon independently of the external supply of combined nitrogen, the nutrient limiting primary production in most ocean regions. Recent investigations indicated that in *Trichodesmium*, a predominant diazotroph in large parts of the oligotrophic oceans, carbon and nitrogen fixation are stimulated by rising pCO₂. In this study we investigate the physiological response of *Nodularia spumigena*, a heterocystous bloom-forming cyanobacterium of the Baltic Sea, to elevated [CO₂] and corresponding acidification as projected for the year 2100. *N. spumigena* reacts with strongly impeded growth and reduced nitrogen fixation. These effects are accompanied by storage of nutrients and significant changes in elemental composition of the cells. Our observation of adverse effects of seawater acidification/carbonation on growth and nitrogen fixation of *N. spumigena* are contrasting previous findings on *Trichodesmium* and eukaryotic marine

²Marine Environmental Laboratories, International Atomic Energy Agency, Monaco.

¹ NIWA, PO Box 56, Dunedin, New Zealand, (k.currie@niwa.co.nz)

² Dept of Chemistry, University of Otago, Dunedin, New Zealand

phytoplankton. Possible explanations for the opposed physiological responses may be found in the different ecological strategies of non-heterocystous *Trichodesmium* and the heterocystous *Nodularia*.

BEHAVIOR OF SUBTROPICAL COASTAL REEF ENVIRONMENTS UNDER RISING ATMOSPHERIC CARBON DIOXIDE AND OCEAN ACIDIFICATION, THE EXAMPLE OF HAWAII AND BERMUDA M12

De Carlo, Eric H.¹, F.T. Mackenzie¹, A.J. Andersson², C.L. Sabine³, and R.A. Feely³

Ocean acidification has potentially significant implications for both carbonate mineral production and dissolution. Here we demonstrate how the response of subtropical reefs to existing conditions can help predict how future higher atmospheric CO₂, lowered seawater pH and carbonate saturation levels will impact marine carbonates. We use results of studies in Hawaii (CRIMP-CO₂) and Bermuda and modeling with the Shallow-water Ocean Carbonate Model (SOCM) to illustrate the response of the subtropical coastal ocean to ocean acidification. Nearly three years of high frequency measurements at CRIMP-CO₂ show that Kaneohe Bay is annually a net source of CO₂ to the atmosphere, mainly due to intense calcification. With CO₂ in bay waters averaging ~450 µatm, if atmospheric CO₂ increases at the rate of 2.1 µatm/yr observed during our study, the bay will switch from being a net source to a net sink of CO₂ within ~30 years. At such time, reefs will increasingly be at risk of decreased calcification rates and particulate carbonate minerals, especially high magnesian calcites, of dissolution. The latter already occursnaturally in Bermuda, where CO₂concentrations reach well above 1000 µatm during the summer in the stratified bottom waters of Devil's Hole, leading to dissolution of phases with an average mol % MgCO₃ of ~16. The SOCM predicts that under these conditions, similar to those predicted in the latter part of this century under the BAU scenario, dissolution of carbonates will exceed production by the year 2150 or sooner, leading to a net loss of carbonate minerals from the world's coastal ocean.

QUANTIFICATION OF pH AND INORGANIC CARBON USING AUTONOMOUS SENSORS T12

DeGrandpre, Mike¹, Sarah Cullison¹, Todd Martz², and Gernot Friederich²

We will present the latest technologies developed in our lab for autonomous quantification of inorganic carbon parameters. An improved version of the Submersible Autonomous Moored Instrument for CO_2 (SAMI- CO_2) that is in the advanced stages of development will be described. A design of a new sensor, SAMI-pH, developed directly from the SAMI- CO_2 technology, is also discussed. Results from recent field studies using SAMI-pH are presented. The sensor has been deployed on Scripps pier, in Monterey Bay (MBARI M0 mooring), in the northeast Pacific, and on a coral reef in Puerto Rico. These studies are characterizing the natural range of pH variability over diurnal to seasonal time scales. Calculation of total dissolved inorganic carbon (DIC) from combined autonomous measurements of the partial pressure of CO_2 (pCO_2) and pH is examined. These preliminary results show that, as expected, the pH- pCO_2 combination is sensitive to errors; however, very good DIC prediction is obtained from both pH and pCO_2 when combined with salinity-derived alkalinity. Our future plans are to combine pH and pCO_2 sensors with an autonomous total alkalinity system currently under development. These combined measurements will make it possible to quantify DIC while simultaneously checking internal consistency of the inorganic carbon parameters.

¹ IFM-GEOMAR, Düsternbrooker Weg 20 24105 Kiel, Germany, (jczerny@ifm-geomar.de)

¹Department of Oceanography, University of Hawaii, Honolulu, HI 96822, USA; edecarlo@soest.hawaii.edu

²Bermuda Institute of Ocean Sciences, 17 Biological Lane, St. George's GE 01 Bermuda

³NOAA PMEL, 7600 Sand Point Way NE, Seattle, WA 98115, USA

² Monterey Bay Aquarium Research Institute, Moss Landing, CA 95039 USA

THE EFFECT OF HIGH CO₂ CONCENTRATIONS ON ENERGY BUDGETS OF ISOLATED GILLS FROM ANTARCTIC NOTOTHENIOIDS M13

Deigweiher, Katrin; Magnus Lucassen, Christian Bock, and Hans-O. Pörtner

Little to no information is available on the sensitivity to hypercapnia of Antarctic fish. Living in a stable environment for Millions of years may make Antarctic fish more sensitive to environmental changes than any other (e.g. temperate or Arctic) species. In this study we investigated the effect of high CO₂ (10.000 ppm) concentrations on energy turnover of fish gills overall as well as on individual energy consuming processes in two Antarctic Notothenioids (Gobionotothen gibberifrons and Notothenia coriiceps). Oxygen consumption of isolated gills was measured under normo- vs. hypercapnic conditions using specific inhibitors to determine the fractional costs of ion regulation, protein and RNA synthesis in the energy budget. The fraction of the three processes was similar in both species. However, absolute costs of all three processes were higher in N. coriiceps than in G. gibberifrons. This may be attributed to the sluggish benthic life style of G. gibberifrons, whereas benthopelagic N. coriiceps actively feed on small fish and krill. Under hypercapnia, gill preparations from G. gibberifrons displayed a drastic rise in energy consumption rates of ion regulation (197%), protein (237%) and RNA synthesis (220%). In contrast, the cost of ion regulation (156%), protein (128%) and RNA synthesis (107%) rose more moderately in N. coriiceps. Evidently, high CO₂ concentrations induce strong increments in the energy demand of cellular processes. Future experiments will show whether these shifts play a role under more realistic CO₂ accumulation scenarios and cause a reduction in energy availability for growth and reproduction.

Dept. Marine Animal Physiology, Alfred-Wegener-Institute for Polar and Marine Research, Am Handelshafen 12, 27570 Bremerhaven, Germany

UNCERTAINTIES IN THE OCEAN CARBONATE SYSTEM AT HIGH P(CO₂) T13 Dickson, Andrew G

The state of the ocean carbonate system is typically described by providing values for two of the four canonical analytical variables: pH, $p(CO_2)$, total alkalinity, and total dissolved inorganic carbon together with the salinity, temperature and pressure so as to allow estimation of the requisite equilibrium constants as well as the total boron concentration. Once this basis set of information is known, it is possible to compute the concentrations of the various carbonate species, and hence infer the remaining analytical variables.

Each of these input variables – including the estimated equilibrium constants – is subject to some uncertainty, resulting in an overall uncertainty that can be ascribed to any individual item of computed information such as the carbonate ion concentration or the pH. In this presentation, I shall discuss how one estimates the likely magnitudes of the various uncertainty terms and infer the overall uncertainty of the various computed information over a range of conditions, including over a range of pressures. In particular, I will examine the previously observed discrepancies for the ratio of the equilibrium constants: K_1/K_2 at high $p(CO_2)$ values, and will indicate the likely consequences of this discrepancy for the accurate prediction of carbonate ion concentrations in systems with high $p(CO_2)$.

Scripps Institution of Oceanography, University of Calfornia, San Diego, 9500 Gilman Drive, La Jolla, California 92093-0244 (adickson@ucsd.edu)

¹ Department of Chemistry, The University of Montana, Missoula, MT 59812 USA (michael.degrandpre@umontana.edu)

IMPACT OF OCEAN ACIDIFICATION ON CALCIFYING MARINE ORGANISMS

M14

Dissard, Delphine, Jelle Bijma Jelle, Regine DaRocha, and Gernot Nehrke

Recently, field and laboratory studies have demonstrated the significant impact of variations in seawater [CO₂] and related changes in carbonate chemistry on both planktic and benthic marine calcifying organisms. (e.g. Bijma et al., 1999; Kleypas et al., 1999; Leclercq, 2000 et al., Riebesell et al., 2000; Zondervan et al., 2001; Delille et al., 2005; Gazeau et al., 2007). For foraminifera a model has been proposed to explain the calcification process (Bentov and Erez, 2005, 2006): Seawater is taken up via endocytosis and transported to the site of calcification. During transport the composition of the solution inside the vesicle is modified (concentrating Ca²⁺ while removing other divalent cations, notably Mg²⁺, and at the same time pumping H⁺ to acidic vesicles, thereby increasing pH and CO₃²⁻ in the "calcification vesicles"). Because the calcification is based on ambient seawater, foraminifers are among the best recorders of paleo-proxies. This also explains why foraminifera are so sensitive to ocean acidification. A similar model has been proposed for corals (Erez, personal com.). The similarity in the ultra-structures of different calcifying organisms also suggests a common ancestral calcification mechanism. We report in more detail on the impact of ocean acidification on planktonic and benthic foraminifera, through controlled laboratory culture experiments. The impact was monitored by weighing shells and using SEM and AFM techniques. The overall aim of this study is to develop/improve a process based understanding of biocalcification in foraminifera which would allow predicting how these calcifiers cope with an acidifying ocean and which consequences this has for the structural properties of their skeletons.

Alfred Wegener Institute for Polar and Marine Research, Germany. (Delphine.Dissard@awi.de, Jelle.Bijma@awi.de)

BORON ISOTOPES IN SCLERACTINIAN CORALS: pH OR TEMPERATURE PROXY? T14 <u>WITHDRAWN</u>

<u>Douville, Eric</u>¹, Pascal Louvat², Jérôme Gaillardet², Guy Cabioch³, Norbert Frank¹, Anne Juillet-Leclerc¹, and Martine Paterne¹

No proxy is perfect! While boron isotopes in tropical corals such as Porites sp. are a powerful tool to accurately reconstruct past temporal changes of the sea surface pH at high resolution, they appear relatively sensible to temperature when they incorporate in North Atlantic deep-sea coral Lophelia pertusa. We present here first results concerning a recent "pH_{seawater} – \Box^{11} B_{coral}" field calibration at seasonal timescale involving modern Porites sp. and surface seawater samples monthly collected from New Caledonia lagoon during 18 months. Obtained data tends to show the high potential of the boron isotopes to reconstruct paleo-pH on seawater scale (SWS) with an accuracy of +/- 0.2 pH-unit including analytical uncertainties. Thus, external reproducibility obtained from boron isotope measurements using MC-ICPMS Neptune was inferior to +/- 0.2 ‰. pH calculations were performed with an isotopic composition of boron for seawater of 39.9 % and an isotopic fractionation factor of 0.981. Such results reinforce the high potential of Porites sp. to quantify Ocean Acidification in tropical latitudes since the beginning of the industrial era. In contrast, preliminary measurements of boron isotopes in deep-sea coral Lophelia pertusa collected between 40°N and 70°N along the European margin of the eastern North Atlantic showed a strong influence of the temperature on the isotopic composition of boron ranging between 26 and 29 ‰. Moreover these relatively high values indicate a strong "vital effect" and their use for paleo-pH reconstruction for intermediate or deep waters require field or experimental calibrations.

¹ LSCE/IPSL, UMR 1572 CNRS-CEA-UVSQ, F-91198 Gif/Yvette, France (eric.douville@lsce.ipsl.fr)

² IRD, B.P. A5, F-98848 Nouméa Cedex, Nouvelle Calédonie

³ IPG-Paris, Laboratoire de Géochimie et de Cosmochimie, Place Jussieu, F-75252 Paris, France.

SURFACE WATER PH MEASUREMENTS AND CARBONATE CHEMISTRY IN THE SUBTROPICAL NORTHEAST ATLANTIC OCEAN M15

<u>Dumousseaud, Cynthia</u>¹, Eric Achterberg¹, Nick Hardman-Mountford², David Hydes¹, Matt Mowlem¹, and Toby Tyrrell¹

In order to get a better understanding of how ocean acidification will affect the plankton community, and particularly calcifying organisms, we need to understand the seasonal biological and physical controls on pH. To observe and consequently predict changes in the ocean, precise measurements of all the carbonate chemistry parameters in the ocean are important. Direct measurement of pH would offer the possibility of making highly detailed observations that can allow precise correlations to be made with the different hydrographic and biogeochemical processes driving changes in pH. Achieving this in practice currently presents a considerable analytical challenge.

A high precision pH measurement system (accuracy and precision of 0.01 pH unit) was tested in the Sub-Tropical Northeast Atlantic during January 2008 aboard the RRS Discovery. High resolution potentiometric pH_F measurements along with partial pressure of CO_2 (pCO₂) were made continuously from the ship's underway supply. Discrete sample were taken for Dissolved Inorganic Carbon (DIC) and Total Alkalinity (TA) measurements. The area studied was under the Saharan dust plume, where atmospheric inputs can be of particular importance on primary productivity in areas where nutrients are limiting. We were able to determine small-scale changes in pH related to response of biology associated with the dust inputs. A comparison of the relative changes in the four carbonate parameters measured will be discussed, in relationship with changes in the physics and biogeochemistry.

CO₂-DRIVEN ACIDIFICATION RADICALLY AFFECTS LARVAL SURVIVAL AND DEVELOPMENT IN MARINE ORGANISMS – III, TUNICATES T15

<u>Dupont, Sam</u>¹, Hélène Auger², Jean-Stéphane Joly², Jean-Marie Bouquet², Daniel Chourrout³, and Michael C Thorndyke³

Tunicates have not been a priority target for research on the consequences of OA, and yet several tunicates are keystone species, aggressive invasives due to their ability to spread rapidly and endanger native taxa, and in some cases a major threat to aquaculture. We have used computerised monitoring and regulation of pH in natural sea water by controlled injection of CO2 to investigate the impact of pH changes predicted by the year 2100 (0.2 – 0.4 pH units) on the long term development of three tunicate species: the sessile species *Ciona intestinalis* and *Ascidiella aspersa*, which can occur at extremely high densities of several thousands of individuals per square meter, and *Oikopleura dioica*, an important component of the pelagic ecosystem bridging the gap between small primary producers and higher consumers. Our results show that tunicates performed better in acidified water for all tested parameters. Notably, tunicates in acidified water grew and developed more rapidly and we observed a significant increase in fecundity. These results will be discussed at the light of potential consequences on future ecosystems.

¹ National Oceanography Centre, European Way, Southampton SO14 3ZH, U.K. (cd6@noc.soton.ac.uk)

² CASIX, Plymouth Marine Laboratory, Plymouth PL1 3DH, U.K.

¹ Dept. Marine Ecology - Kristineberg, University of Gothenburg, Kristineberg 566, 45034 Fiskebäckskil, Sweden (sam.dupont@marecol.gu.se)

² Institut Fessard, CNRS - 91198, Gif-sur-Yvette, France

³ Sars International Centre for Marine Molecular Biology - 5008 Bergen, Norway

A NUMERICAL STUDY OF TRANSPORT AND SPREADING OF GASES FROM GASSEEPAGE THROUGH THE SEAFLOOR M16

Enstad, Lars Inge^{1,2}, Peter M. Haugan², and Guttorm Alendal^{1,3}

Gas seeps through the seafloor at several locations around the world oceans. These seeps are observed as both CO₂ rich liquid/gas in volcanic areas such as Mid Ocean Ridges, and as hydrocarbon gases, consisting mainly of CH₄, in hydrocarbon rich areas such as the North Sea. The release of CO₂ through the ocean bottom is a contribution to the global CO₂ budget. These events are interesting from a phenomenological perspective and have been studied through observations and modeling the last decades. In order to describe the role of such sources in ocean biogeochemical cycles, it is necessary to understand how the gas is dissolved and spreads in plumes above the leakage. In this investigation we combine modeling efforts on the dissolution and spreading from the leakage point and compare with in situ data.

In this specific study a modeling tool has been developed where the MIT general circulation model (http://mitgcm.org/) has been coupled with an improved version of a single bubble/droplet model. The MITgcm model has previously been modified to make it capable of simulating transport and spreading of CO₂ as a dynamically active tracer. The vertical mixing of CO₂ is parameterized state-of-the-art models using the Ocean Turbulence employing General (http://www.gotm.net/). The coupled droplet/bubble-OGCM has been compared with the existing single-droplet model and also plume models to verify the implementation. At the current stage the gas flux through the seafloor is given at a fixed rate, with a statistical distribution to set the bubble/droplet sizes.

To check the model capabilities in a natural environment the model will be applied to an area where there exist in-situ data for the environmental parameters and for the plume spreading. We hope to be able to present comparison using different parameters of these data with model results at the conference.

THE PROCESSES INFLUENCING THE DISTRIBUTION OF INTERTIDAL COMMUNITIES IN A HIGH ${\rm CO}_2$ OCEAN T16

Findlay, H.S.¹, M.A. Kendall¹, J.I. Spicer², S. Widdicombe¹, and C. Turley¹

The barnacle *Semibalanus balanoides* is a major space occupier on rocky shores in Northern Europe and hence changes in its population ecology can have a broad influence on other species. Despite its ecological importance, the impact of a warmer, more acid ocean on this species, and hence on intertidal communities, remains unclear. In this paper we demonstrate how temperature and carbon dioxide interact to affect *S. balanoides* egg development, nauplii development and cyprid development. Changes in abundance and viability of the early life stages will impact the supply of larvae arriving in the intertidal and post settlement mortality will determine the number of individuals reaching reproductive age. Laboratory experiments indicated that changes in pH and temperature slowed the metamorphosis of cyprids thereby increasing their exposure to desiccation. Nevertheless, increased temperature and CO₂ concentration had greatest impact on smaller individuals prior to metamorphosis; with poor survival being linked to slow growth and ability to calcify. The paper also also discusses how this experimental data can be used to develop models for the prediction of changes in population distributions as well as aspects such as secondary production and calcium carbonate production.

¹Plymouth Marine Laboratory, Prospect Place, West Hoe, Plymouth, PL1 3DH, U.K. (hefi@pml.ac.uk)

¹ Bergen Center for Computational Science, Unifob, Thormøhlensgate 55, 5008 Bergen email: lars.inge.enstad@bccs.uib.no

² Geophysical Institute, University of Bergen, Allegaten 70, 5007 Bergen

³ Department of Mathematics, University of Bergen, Johannes Brunsgate 12, 5008 Bergen

THE IMPACT OF MODEL RESOLUTION ON PROJECTIONS OF FUTURE OCEAN ACIDIFICATION M17

Gangstø, Reidun^{1,2}, Fortunat Joos^{2,3}, and Marion Gehlen¹

Ocean acidification with the associated shift of the calcite and aragonite saturation horizons is likely to have a negative impact on pelagic organisms producing calcite and aragonite shells, such as coccolithophores, foraminiferas and pteropods. In order to quantify these effects, comprehensive and thorough model studies of both the changing calcite and aragonite saturation state and its effect on calcite and aragonite production and dissolution are essential.

We have used the pelagic biogeochemical ecosystem model PISCES in combination with two dynamical models of different resolutions. The PISCES model simulates the marine biological productivity and describes the biogeochemical cycles of carbon and main nutrients. It distinguishes two phytoplankton and two zooplankton size classes. The production of calcite is assigned to nanophytoplankton as a function of temperature, saturation state, nutrient and light availability, whereas the production of aragonite is assigned to mesozooplankton as a function of saturation state. The PISCES model is originally connected to the global dynamical NEMO/OPA8.2 model which is of relatively high resolution. We have additionally coupled PISCES to the global dynamical low spatial resolution Bern3D model. Both the OPA/PISCES model and the new Bern3D/PISCES model were run until equilibrium state was reached and the models were forced with increasing atmospheric CO₂ concentrations following the SRESA2 scenario. Here we show the effect from using two dynamical models of different resolutions on the projected ocean acidification in addition to presenting potential changes in the pelagic calcite and aragonite budget with increasing concentrations of atmospheric CO₂.

EUROPEAN PROJECT ON OCEAN ACIDIFICATION (EPOCA) T17

Gattuso, J.-P., L. Hansson, and the EPOCA Consortium

The European Project on Ocean Acidification (EPOCA) was launched in May 2008 with the overall goal to advance our understanding of the biological, ecological, biogeochemical, and societal implications of ocean acidification. Its consortium includes 105 principal investigators from 27 institutes. The budget of this 4 year long project is 16,5 M€, including 6,5 M€ from the European Union. The research efforts of EPOCA are divided into four different themes. First, EPOCA will focus on past and present spatiotemporal changes in ocean chemistry and biogeography of key marine organisms. Second, EPOCA will quantify impacts of ocean acidification on marine organisms and ecosystems. Molecular, physiological and ecological approaches will be used to study climaterelevant biogeochemical processes, including calcification, primary production and nitrogen fixation. Laboratory and field perturbation experiments will focus on key organisms in terms of their ecological, biogeochemical, or socioeconomic importance. Third, EPOCA will improve biogeochemical, sediment, and coupled ocean-climate models to better account for how ocean acidification will affect ocean biogeochemistry and ecosystems. Special attention will be paid to feedbacks of physiological changes on the carbon, nitrogen, iron, and sulfur cycles and how these changes will affect and be affected by future climate change. Finally, EPOCA will evaluate uncertainties, risks and thresholds (tipping points) related to ocean acidification at molecular, cellular and organismal levels from local to global scales. It will also assess the decrease in CO₂ emissions required to avoid these thresholds and describe the change to the marine environment and Earth system, should these emissions be exceeded.

¹ Laboratoire des Sciences du Climat et de L'Environnement (LSCE), Gif-sur-Yvette, France

² Climate and Environmental Physics, Physics Institute, University of Bern, Switzerland (gangsto@climate.unibe.ch)

³ Oeschger Centre for Climate Change Research, University of Bern, Switzerland

EPOCA Project Office: Laboratoire d'Océanographie, BP 28, F-06234 Villefranche-sur-mer Cedex, France, (coord-ocean-acidification@obs-vlfr.fr), http://epoca-project.eu/

EFFECT OF SEAWATER ACIDIFICATION ON THE GROWTH OF MYTILUS EDULIS M18

<u>Gazeau, Frédéric</u>¹, Aimé Roger Nzigou¹, Hannah Wood², Jean-Pierre Gattuso³, Jack Middelburg¹, and Carlo Heip¹

The Intergovernmental Panel on Climate Change (IPCC) predicts atmospheric CO₂ partial pressure (pCO₂) ranging from 490 to 1,250 ppmv in 2100, depending on the socio-economic scenario considered. Because one third of anthropogenic CO₂ emissions has been stored in the oceans, ocean pH has already declined by 0.1 unit compared with pre-industrial values and is predicted to decrease by another 0.4 unit by the end of the century. Seawater acidification lowers the concentration of carbonate ions, one of the building blocks of calcium carbonate (CaCO₃), which can in turn greatly affects the ability of calcifying organisms to precipitate CaCO₃. Few studies have investigated the detrimental effect of acidic waters on bivalves and none investigated the response of calcification to pCO₂ levels within the range of values projected by IPCC. Here, we report experimental data showing that calcification and growth of mussels (Mytilus edulis) is significantly impacted by a decrease of seawater pH. In the laboratory, mussels were incubated during 4 months at two different pCO₂: 400 and 800 ppm. Calcification rates estimated using the alkalinity anomaly method and shell length vs. shell weight relationships showed a significant decrease of calcareous production between the 400 and the 800 ppm treatment. A shift in shell mineralogy has also been evidenced as aragonite/calcite ratios decreased significantly with increased pCO₂. Finally, an important decrease of mussel health determined using the Neutral Red Retention technique has been shown at 800 ppm, suggesting elevated levels of calcium ions in the haemolymph, generated by dissolution of the shells.

² Plymouth Marine Laboratory, Plymouth, U.K.

21ST-CENTURY CHANGES IN BOTTOM-WATER PH IN THE HIGH-LATITUDE NORTH ATLANTIC. T18

Gehlen, M.¹, L. Bopp¹, P. Cadule¹, D. Swingedouw², and J. C. Orr³

Data-based estimates indicate that anthropogenic CO₂ has already penetrated deep into water masses of the abyssal high-latitude North Atlantic. A suite of models hindcasting similar present-day patterns project that if atmospheric CO₂ continues to increase following the IPCC IS92a (business-as-usual) scenario, these same water masses will reach levels of anthropogenic DIC by 2100 exceeding 100 umol kg⁻¹. This is twice the present-day surface concentration. We translated these perturbations into corresponding reductions in pH of bottom waters and found that they exceed 0.2 pH units in most bottom waters of the North Atlantic situated poleward of 50°N. Reductions in pH beyond this threshold are considered by environmental agencies to be dangerous in natural waters, and have been reported to be detrimental to deep-sea benthic organisms. Moreover, these pH reductions are much larger than the amplitude of natural variability of pH in North Atlantic bottom waters. To study how effects of climate change may alter projected changes in bottom-water pH, e.g., by slowing Atlantic Meridional Overturning, we exploited simulations from a coupled climate-carbon cycle model and a sensitivity test to isolate effects from melting of land ice. Generally, 21st-century climate change was found to render bottom-water pH reductions somewhat less severe, especially when considering effects from land ice melt. Yet in all simulations, with or without climate change, there are large reductions in pH in bottom waters of the North Atlantic. These will expose sediments and benthic organisms to changes in conditions during the 21st century that are much larger than any variability that they have experienced during at least the last 420,000 years.

¹ Netherlands Institute of Ecology, Yerseke, The Netherlands (f.gazeau@nioo.knaw.nl)

³ Laboratoire d'Océanographie de Villefranche, Villefranche-sur-Mer, France

EFFECT OF SEAWATER ACIDIFICATION ON EPIBIONT-BEARING CIDAROID SPINES M19

Guibourt, Virginie, Ana I. Catarino, and Philippe Dubois

Due to their well-developped high-magnesium calcite endoskeleton, echinoderms will particularly suffer from seawater acidification. Among them, cidaroid sea urchins could be especially affected since, in this group, fully-grown spines are no more covered by an epidermis, leaving the skeleton in direct contact with seawater. These « naked » spines are usually heavily colonized by epibionts making these sea urchins « islands of biodiversity ». This is particularly well evidenced in the Southern Ocean where the presence of cidaroids and their epibionts significantly increase biodiversity of the muddy substrates on which they live. Most models of the ocean-carbon cycle predict that the shallowing of the calcium carbonate saturation horizons due to increasing anthropogenic CO₂ emissions will be particularly important in the Southern Ocean. So, effects of acidification on cidaroid spines are particularly relevant to the ecology of this region.

In the present study, we assessed the impact of magnesium-calcite saturation level on cidaroid spines of both experimental and field specimens. In experimental conditions, spines were incubated for up to three weeks in CO₂ enriched seawater of controlled pH (7.2, 7.6 and 8.2) and temperature. Total alkalinity was measured and saturation level calculated. The spine immerged weight was measured, morphological evidence of etching was monitored by scanning electron microscopy and cover of epibionts was evaluated. Dried field specimens from different depths below and above the current saturation horizon were obtained from museum collections. Morphological evidence of etching and magnesium concentration in the skeleton were determined.

Laboratoire de Biologie marine CP 160/15, Université Libre de Bruxelles, 50 av Roosevelt, B-1050 Bruxelles, Belgium (phdubois@ulb.ac.be)

GROWTH, CALCIFICATION AND ACID-BASE REGULATION IN THE CEPHALOPOD SEPIA OFFICINALIS UNDER ELEVATED ${\rm CO}_2$ T19

Gutowska, Magdalena A.¹, Frank Melzner², Martina Langenbuch¹, Franz J. Sartoris¹, and Hans O. Pörtner¹

Changes in seawater pH and carbonate chemistry associated with ocean acidification negatively influence calcification in all marine invertebrates studied to date. Our experimental work, with the cephalopod mollusc *Sepia officinalis*, is the first to show that in some invertebrates, neither calcification nor somatic growth decrease under low seawater pH and low Ω_{arag} . During a six-week period, juvenile *S. officinalis* maintained under both ~4,000 and ~6,000ppm CO₂ grew at the same rate (4% body mass daily) and gross growth efficiency as control animals, increasing the mass of their calcified internal shell by >500%. Closer examination of the tissue and blood acid-base status of the animals revealed that during long-term exposure they had fully compensated the initial acidification and maintained elevated bicarbonate levels in their blood. Accumulated bicarbonate did not result from internal shell dissolution as in other molluscs, extracellular [Ca²⁺] was maintained at control levels. On the contrary, high CO₂ (6,000 ppm) animals were able to deposit significantly higher amounts of CaCO₃ in their shells and also were characterized by altered shell morphology. SEM images revealed strong increases in the number of shell-chambers constructed by high-CO₂ animals. Whether observed changes in shell morphology under high-CO₂ conditions constrain the ecological fitness of the animal in any way needs to be investigated. We conclude that cephalopods possess a

¹ Laboratoire des Sciences du Climat et de l'Environnement (LSCE/IPSL), CEA/CNRS/UVSQ, CE Saclay, Orme des Merisiers, 91191 Gif-sur-Yvette

² European Centre for Research and Advanced Training in Scientific Computation (CERFACS), 42 Avenue Gaspard Coriolis, 31057 Toulouse

³ Marine Environment Laboratories (MEL-IAEA), 4, Quai Antoine 1^{er}, MC-98000 Monaco

certain level of pre-adaptation to long-term increments in carbon dioxide levels, as indicated by their efficient pH compensation, maintained growth and calcification performance.

BIOGEOCHEMICAL APPROACH OF ENVIRONMENTAL ASSESSMENT FOR CO2 OCEAN STORAGE M20

Harada, Koh, Nobuo Tsurushima, Masahiro Suzumura, and Namiha Yamada

Ocean should be a good candidate for storage of CO₂ collected from large sources because it can receive a large amount of CO₂ without significant change of pH and DIC due to its large water mass. However, significant changes in chemical properties could occur near release points of CO₂. Storage of CO₂ into geological formation under seabed is also considered and it has a possibility of leakage of CO₂ into bottom water. If the significant changes in chemical properties occur near the release and/or leakage points, not only marine organisms but also biogeochemical processes can be affected. To understand effects onto the biogeochemical processes including dissolution and degradation of inorganic and organic particles, we studied how low pH and high CO₂ can effect onto dissolution of calcium carbonate and also activities of microbial processes.

Laboratory experiments were conducted under high pressure and high pCO₂ condition to understand change of the dissolution rate of calcium carbonate. As a result, rapid dissolution was observed at above 5000 ppm of pCO₂. The initial dissolution rate was well correlated with initial concentration of dissolved inorganic carbon. Dissolution of calcium carbonate increased dissolved inorganic carbon and total alkalinity in the seawater. Then, degrees of undersaturation and dissolution rate of calcite in the seawater were decreased with time. Dissolution rates normalized with apparent surface area were well fit to the rate law in the empirical kinetic showed in previous studies.

We will also show result of laboratory experiment for bacteria growth in our poster.

Research Institute for Environmental Management Technology, National Institute of Advanced Industrial Sceinece and Technology, 16-1 Onogawa, Tsukuba, Ibaraki 305-8569, Japan (koh.harada@ni.aist.go.jp)

FUTURE ECOSYSTEM CHANGES PROJECTED BY A 3-D HIGH-RESOLUTION ECOSYSTEM MODEL T20 <u>WITHDRAWN</u>

Hashioka, Taketo^{1,2}, Takashi T. Sakamoto¹, Takeshi Okunishi³, Hiroshi Sumata⁴, and <u>Yasuhiro Yamanaka</u>^{1,2,4}

In recent years impacts of global warming on physical environment have been projected through some scenario experiments using climate models (CO-AGCMs). As a challenge of modelling responses to global warming on marine ecosystem, we developed a 3-D high-resolution ecosystem model, COCO-NEMURO, with off-line calculation method which can directly use projected results of CO-AGCMs as a physical field of the ecosystem model. The COCO-NEMURO consists of PICES NEMURO (North Pacific Ecosystem Model for Understanding Regional Oceanography) coupled with COCO (CCSR Ocean Component Model) which has a horizontal resolution of 1/4 by 1/6 degrees. As a first step, we applied this model to the western North Pacific, and conducted a global warming experiment using physical fields from a high-resolution climate model (the CCSR/NIES/FRCGC CO-AGCM which contributed to the IPCC-AR4, and will contribute to the AR5). The experiment is conducted following a scenario in which atmospheric CO₂ concentration increases by 1% per year. Our model well reproduced the seasonal and regional variations of lower-trophic level ecosystem associated with meso-scale features in the present-day simulation. Under the global warming condition, it is interesting that our model projected the increase in Chl-a concentration during spring bloom by 10 to 20 % in the subarctic region even though annually averaged Chl-a concentration decreases. Since

¹Alfred-Wegener-Institute for Polar and Marine Research, Bremerhaven, Germany (Magdalena.Gutowska@awi.de)

²Leibniz Institute for Marine Sciences, IFM-GEOMAR, Kiel, Germany.

these changes would affect the higher-trophic level ecosystem, we have been also developing an integrated marine ecosystem model explicitly representing linkages between the lower-trophic level ecosystem and major pelagic fishes based on COCO-NEMURO. We present some highlights in our first stage results.

¹ Frontier Research Center for Global Change (FRCGC) / Japan Agency for Marine-Earth Science and Technology (JAMSTEC), 3173-25, Showa-machi, Kanazawaku-ku, Yokohama, Japan (galapen@ees.hokudai.ac.jp)

² Core Research for Evolutional Science and Technology (CREST) / Japan Science and Technology Agency (JST), 4-1-8, Honcho, Kawaguchi-shi, Japan

³ Tohoku National Fisheries Research Institute, Fisheries Research Agency (FRA), 3-27-5, Shinhamacho, Shiogama, Miyagi 985-0001, Japan.

⁴ Hokkaido University, Faculty of Environmental Earth Science, N10W5, Kita-ku, Sapporo, Japan

OBSERVATIONS OF ACIDIFICATION IN THE WEDDELL SEA ON A DECADAL TIME SCALE M21

Hauck, Judith¹, Mario Hoppema¹, Christoph Völker¹, Richard Bellerby², and Dieter Wolf-Gladrow¹

The amount of anthropogenic CO_2 (C^{ant}) that entered the Weddell Sea between 1992 and 2008 is assessed using extended multiple linear regression. Two multiple linear regressions for total dissolved inorganic carbon (C_T) were conducted independently for two data sets from 1992 and 2008 based on potential temperature, salinity and oxygen. Subtracting these two relationships leads to an estimate of C^{ant} accumulated in the considered time span (ΔC_T) assuming that the relations between input parameters and C_T do not change with time.

Invasion of C^{ant} appears to have occurred in agreement with the general knowledge of the circulation indeed. In the Antarctic Circumpolar Current and in the Warm Deep Water ΔC_T values are close to zero, whereas values as high as 10 μ mol kg⁻¹ are observed in the surface layer. ΔC_T concentrations at the surface vary with latitude between 1 and 15 μ mol kg⁻¹. Also the intrusion of C^{ant} into deep layers was demonstrated, near the sea floor ΔC_T yields 5–6 μ mol kg⁻¹.

The invasion of C^{ant} provokes a shift in the equilibria of the carbonate system, resulting in acidification and depletion of CO_3^{2-} . A detailed account of the changes that occurred between 1992 and 2008 will be given. The mean decrease of pH in the upper 200 m layer is 0.015.

Further effects are decrease of calcite and aragonite saturation levels. In contrast to other studies, results of this analysis suggest that aragonite undersaturation of surface waters in the Weddell Sea won't be reached before the year 2150.

THE UTILIZATION OF PHYSIOLOGICAL MEASUREMENTS TO PREDICT FUTURE 'WINNERS' IN A HIGH-CO $_2$ WORLD – A CASE STUDY USING THE MODEL AMPHIPOD GAMMARUS LOCUSTAi T21

Hauton, Chris, Toby Tyrrell, and John Williams

It is almost universally accepted that the current atmospheric CO₂ concentration of ca. 380 ppmv is set to rise by an estimated 1% y⁻¹ over the next few decades. However, it currently remains difficult to predict the future impacts of acidification, especially in coastal and estuarine species which may have evolved to accommodate episodic low pH environments.

We report an investigation of the effects of increases in pCO₂ on the growth and molecular physiology of the neritic amphipod *Gammarus locusta*, which has a cosmopolitan distribution in estuaries. Amphipods were reared from juvenile to mature adult (~ 28 days) in laboratory mesocosms at three

¹Alfred-Wegener-Institute for Polar and Marine Research, Postbox 12 01 61, D-27515 Bremerhaven, Germany (judith.hauck@awi.de)

² Bjerknes Centre for Climate Research, University of Bergen. Allégaten 70, 5007 Bergen, Norway

different pCO₂ concentrations, pCO₂ was regulated by computer-controlled addition of 100% CO₂ which maintained pH set points of nominally pH 8.1 (control) and pH 7.8 and 7.6 (experimental).

Growth rate was estimated from weekly measures of body length and converted to AFDW. At sexual maturity the amphipods were sacrificed and assayed for changes in expression of the genes coding for a heat shock protein and the metabolic enzyme glyceraldehyde-3-phosphate dehydrogenase.

The presented data show that this species of amphipod is not significantly impacted by a decrease in sea water pH of up to 0.5 units. These data highlight the fact that in a future 'High CO₂ World' there will be winners as well as losers, which may lead to shifts in the biogeographic distribution of many species. Future experiments modeling the effects of increased pCO₂ on the reproductive success and other trans-generational impacts will be discussed.

School of Ocean and Earth Science, University of Southampton, National Oceanography Centre, European Way, Southampton, Hants, SO14 3ZH, U.K. ch10@noc.soton.ac.uk

CO2-DRIVEN ACIDIFICATION RADICALLY AFFECTS DEVELOPMENT AND LARVAL SURVIVAL IN MARINE ORGANISMS – I. IMPACT ON FERTILIZATION M22

Havenhand, Jon¹, Michael C Thorndyke², Jane E Williamson³, and Sam Dupont²

CO₂-induced ocean acidification threatens the viability of keystone calcifying taxa such as corals, coccolithophores, and pelagic molluscs. Research to date has focussed on the adult stages of calcifying taxa, using gross pH changes relevant for the years 2200 - 2400. We investigated the consequences of exposure to CO₂-induced acidification by -0.4 pH units (the upper limits of predictions for the year 2100, IPCC AR4 2007) for gametes and larvae of keystone marine species. We found statistically significant reductions in sperm swimming performance, fertilization success, and post-metamorphic juvenile survival in acidified treatments. We discuss the implications of these findings for fertilization success and development of the larvae of both non-calcified and calcified taxa, and for the population viability of marine invertebrates.

GLACIAL LESSONS FOR HIGH CO₂ – LIMITATIONS AND CHANCES **T22** Heinze, Christoph

Glacial atmospheric CO₂ partial pressures were about 200 uatm lower than today and 100 uatm lower than at the preindustrial. It is tempting to use the glacial low CO₂ world as a reverse paleo-analog to a high CO₂ world. For the ocean, the last climatic cycle is very well documented by marine sediment core data. However, the glacial low CO₂ world was probably created through an internal redistribution of "already circulating carbon" between the Earth system reservoirs, rather than through extracting or adding carbon to or from an external reservoir (fossil fuel resources). Apart from this fundamental difference, several questions with relevance to the high CO₂ world may be answered through glacial evidence: (1) Was the CaCO₃:POC rain ratio enhanced under higher pH conditions at the sea surface? (2) What were the consequences of high DIC concentrations in glacial deep waters? (3) Why seem both the physical and the biological CO₂ feedback to act positively for glacial-interglacial changes? For addressing these questions, the method of synthetic sediment cores was developed. We demonstrate how it can be used to combine biogeochemical ocean general circulation models with sediment and ice core data to inversely estimate the glacial-interglacial changes in governing carbon cycle parameters. The method is the foundation for a systematic data assimilation of the marine paleo-

¹ Dept. Marine Ecology - Tjärnö, Göteborg University, 45296 Strömstad, Sweden email: jon.havenhand@marecol.gu.se

² Sven Lovén Centre for Marine Sciences, Göteborg University, 566 Kristineberg, 45034 Fiskebäckskil, Sweden

³ Marine Ecology Group, Biological Sciences, Macquarie University, NSW 2109, Australia

record into Earth system models for calibrating their sensitivity to internal and external forcings. It opens the possibility to include all our knowledge of the past in models for predicting an as yet uncertain future.

University of Bergen, Geophysical Institute & Bjerknes Centre for Climate Research, Allégaten 70, N-5007 Bergen, Norway (christoph.heinze@gfi.uib.no).

HYPERCAPNIC STRESS IN EARLY LIFE STAGES OF THE SQUID LOLIGO VULGARIS DUE TO ELEVATED ENVIRONMENTAL CO₂ M23

Hu, Marian Y.-A., Andrea Y. Frommel, Catriona Clemmesen, and Frank Melzner

Cephalopods are high performance organisms and key predators in marine food webs. The effects of rising CO₂ and resulting changes in the ocean's carbonate system on these animals are yet poorly understood. In this study, we test the impact of elevated CO₂ concentrations on the development and performance of early life stages of cephalopods. Loligo vulgaris eggs were incubated at 15°C in sea water equilibrated with three different CO₂-air mixtures (380ppm, 1,400ppm or 4,000ppm CO₂ fraction). Under high CO₂ concentrations (4,000ppm), stage 26 embryos exhibit significantly shorter dorsal mantle lengths and slower developmental rates when compared to larvae incubated at lower CO₂ levels (380ppm and 1,400ppm). After hatch, somatic growth rates will be compared to biochemical indicators of growth and condition. Protein and lipid concentrations, as well as RNA/DNA ratios will serve as measures for energy stores and growth potential, respectively. The energetic investigations are complemented by the study of ion regulatory structures, which are highly challenged by acid-base disturbances. Thus, special emphasis is given to the development of branchial epithelia, by immunohistochemical methods to follow the structural differentiation of gill tissue as well as the occurrence and localization of the marker protein Na⁺/K⁺-ATPase. Additional information will be provided by the screening of the ion regulatory transcriptome (e.g. Na⁺/H⁺-exchangers, HCO₃⁻ transporters, Na⁺/K⁺-ATPase etc.) of whole embryos and various tissues such as gill, branchial heart and skin.

Leibniz Institute of Marine Sciences (IfM-GEOMAR), Hohenbergstr. 2, 24105 Kiel, Germany, mhu@ifm-geomar.de

AN OBSERVATIONAL AND MODELLING STUDY TO DEFINE THE CURRENT pH STATUS OF U.K. WATERS T23

<u>Hydes, David</u>¹, Eric Achterberg¹, Dorothee Bakker², Jerry Blackford³, Nick Hardman-Mountford³, Matt Mowlem¹, Ute Schuster², Carol Turley³ Toby Tyrrell¹ and Andew Watson²

Although a number of serious potential impacts of marine acidification are recognised, little information exists on what present conditions are particularly in the complex environment of European shelf seas. The U.K. Government has funded two years of "baseline" observations. Relatively large natural changes occur in pH through a seasonal cycle. The measurement programme will determine the scale of these changes in a range of areas from eutrophic estuarine to open ocean. New measurements will be linked with on-going monitoring which will provide the information required to put the observations into their hydrodynamic and biogeochemical context. Existing time series data will be used along with the new measurements to quantify the likely degree of variation in pH over longer periods. A data set will be established of the new data and metadata describing its collection and validation. The carbonate system will be studied through measurements of discrete samples to determine DIC and Total Alkalinity, with continuous determination of pCO₂ on related cruises and ships of opportunity and direct measurements of pH. The newly available data will be used to evaluate and improve the ERSEM-POLCOMS full-shelf dynamic simulation model.

¹ National Oceanography Centre, Southampton SO14 3ZH, U.K. (djh@noc.soton.ac.uk)

² University of East Anglia, Norwich, NR4 7TJ, U.K.

³ Plymouth Marine Laboratory, Plymouth PL1 3DH, U.K.

A TOOL FOR EARLY DETECTION OF GLOBAL-SCALE CHANGES IN MARINE CALCIFICATION M24

<u>Ilyina, Tatjana</u>¹, Richard E. Zeebe¹, Ernst Maier-Reimer², and Christoph Heinze³

Ocean acidification is likely to impact calcification rates in many pelagic organisms, which may in turn cause significant changes in marine ecosystem structure. We examine effects of changes in marine CaCO₃ production on total alkalinity (TA) in the ocean using the global biogeochemical ocean model HAMOCC and a variety of future calcification scenarios. The model integrations start at a preindustrial steady state in the year 1800 and run until the year 2300 forced with anthropogenic CO₂ emissions. Calculated trends in TA are evaluated taking into account the natural variability in ocean carbonate chemistry, as derived from repeat hydrographic transects. Using a synthesis of modeling and observational data, we address the following questions. (1) At what time does ocean acidification lead to a decline in marine calcification on a global scale that is detectable in the data? (2) Assuming different calcification scenarios, does the data presently available already allow discerning significant trends, given natural variability and data uncertainties? (3) If so, what is the magnitude and time scale of the decline? Our long-term goal is to help identifying target regions for future ocean chemistry surveys that are critical for early detection and determination of the exact magnitude of large-scale acidification effects from field data.

EFFECTS OF ELEVATED CO₂ CONCENTRATIONS ON P-UTILIZATION OF THE DIAZOTROPHIC CYANOBACTERIUM *NODULARIA SPUMIGENA* AND A NATURAL SUMMER PHYTOPLANKTON COMMUNITY IN THE BALTIC SEA T24

Isensee, Kirsten¹, Henning Johansen¹, Maren Voss¹, Ulf Riebesell² and Monika Nausch¹

The influence of elevated pCO₂ concentrations on cyanobacteria of the Baltic Sea was examined in two different approaches:

- 1. In free drifting mesocosms (60 m³) a gradient of different pCO $_2$ levels was adjusted by acidification with HCl. P-utilization ([33 P]PO $_4$ uptake) positively correlated with increasing pCO $_2$ for small coccoid cyanobacteria. In contrast, cyanobacteria in the size fraction >10 μ m showed less P uptake under pCO $_2$ levels above 1000 ppm. Measurements of the inorganic and organic phosphorus pool (DIP, DOP) indicated no significant differences in consumption for the bulk phytoplankton along the CO $_2$ gradient.
- 2. In laboratory experiments with colony forming *Nodularia spumigena*, a representative Baltic Sea cyanobacterium, we investigated the kinetic response of P-utilization of DIP and DOP. We incubated *N. spumigena* under defined pCO₂ values which we obtained by directly injecting CO₂ gas to reach 450 ppm, 600 ppm and 900 ppm in 0.2 µm filtered natural seawater. DIP and DOP were added and their uptake and nitrogen fixation rates measured. The results suggested a lower P-utilization up to 64 % under present nutrient conditions and high pCO₂ levels, while additional DIP tends to favour the growth under high pCO₂. At pCO₂ of 900 ppm we detected 75 % higher P-utilization compared to present day conditions. In contrast, the amendment of DOP does not stimulate the P-utilization of *N. spumigena* under high pCO₂ levels.

Cyanobacteria with heterocysts and unicellular species may respond differently to elevated CO₂ concentrations which may results in possible changes in species composition with rising atmospheric pCO₂.

¹School of Ocean and Earth Science and Technology, Department of Oceanography, University of Hawaii, 1000 Pope Road, Honolulu, HI 96822, USA. (ilyina@soest.hawaii.edu)

²Max Planck Institute for Meteorology, Bundesstr. 53, 20146 Hamburg, Germany.

³University of Bergen, Geophysical Institute and Bjerknes Centre for Climate Research, Norway.

¹Leibniz Institute for Baltic Sea Research (IOW), Department of Biological Oceanography, Seestrasse 15, D-18119 Rostock, Germany (kirsten.isensee@io-warnemuende.de)

²Leibniz Institute of Marine Sciences (IFM-GEOMAR), Düsternbrooker Weg 20, D-24105 Kiel, Germany

IMPACT OF CLIMATE VARIABILITY ON SURFACE OCEAN CO₂ IN THE PACIFIC SIMULATED IN A 3-DIMENSIONAL MODEL M25

Ishida, Akio 1,2 , Maki N. Aita 1,2 and Yasuhiro Yamanaka 1,2,3

Interannual to decadal variability of carbon cycle is diagnosed with a 3-dimensional physicalbiogeochemical model to improve our understanding of physical and biological impact on the carbon cycle in the Pacific Ocean. We have performed the simulations with two boundary conditions for atmospheric pCO₂: one using the historical increase in atmospheric pCO₂ from year 1783 to 2002 (historical run), another with a constant pre-anthropogenic concentration of 278 ppmv (control run) in order to quantify the natural and anthropogenic carbon cycle. The modeled surface ocean at the Hawaiian Ocean Time-series (HOT) shows a long tern shift in carbonate equilibrium to lower pH and lower saturation states of the carbonate mineral aragonite, which are consistent with the observation. The model simulates two dominant climate variations; Pacific Decadal Oscillation (PDO) and El Niño/Southern Oscillation (ENSO). In the central North Pacific, primary production and biomass increase after the climate shift during the mid 1970s, and CO₂ flux also exhibits interannual-decadal variability. The variations of natural and anthropogenic CO₂ flux are in phase, i.e., both increase and decreases in the Central Pacific in interannual to decadal scale. The variations of natural and anthropogenic CO2 flux are out of phase in the eastern Pacific, i.e., air-to-sea flux of anthropogenic CO₂ decreases when natural CO₂ flux increases and vice versa. This is explained with physical conditions such as upwelling and thermocline variability associated with El Niño and La Niña. Interannual variability of surface saturation states of the carbonate minerals associated with the climate variability is presented.

¹Frontier Research Center for Global Change, Japan Agency for Marine-Earth Science and Technology (JAMSTEC), 3173-25, Showa-machi, Kanazawa-ku, Yokohama, 236-0001, Japan. (ishidaa@jamstec.go.jp)

²Core Research for Evolutional Science and Technology (CREST), Japan Science and Technology Agency (JST), Kawaguchi, 332-0012, Japan.

³Graduate School of Environmental Earth Science, Hokkaido University, N10W5, Kita-ku, Sapporo, 060-0810, Japan.

IN SITU ENCLOSURE EXPERIMENT DEVICE FOR ASSESSING DEEP-SEA ECOSYSTEMS WITH HIGH CO₂ CONCENTRATIONS T25

Ishida, Hiroshi^{1,2}, Yuji Watanabe^{1,2}, Michimasa Magi² and Yoshihisa Shirayama³

It has been proposed to inject CO₂ into the ocean as a potentially effective method to mitigate global warming. Various methods such as seabed direct injection and dilution/dissolution techniques have been proposed, and their feasibility studies being carried out. However, the information on the impact of high CO₂ concentrations on deep ocean ecosystems is insufficient. In order to study the influence of CO₂ sequestration on the deep-sea ecosystem, it is necessary to investigate the ecosystem not only at the species level but also at the community level. To asses the influence of high CO₂ concentrations on deep-sea organisms precisely, it is necessary to carry out *in situ* experiments in deep-sea.

We here developed two types of experimental devices for evaluating on deep-sea ecosystems with elevated CO_2 concentration. One is a free fall type benthic chamber system on deep-sea benthic communities. The other is a water enclosure experiment system (pelagic chamber) in order to do the *in situ* experiment on plankton communities at 1,000 - 2,500 m in depth. Experiments with the benthic chamber system with elevated CO_2 concentrations (averages of $20,000\mu$ atm, $5,000\mu$ atm,

2,000µatm and control) were carried out in the Kumano Trough. The pelagic chamber system is under development and will consist of three chambers (250 L x 3), CO₂ control and water sampling apparatuses with several kinds of sensors. Results of the experiments using these systems and future planning of *in situ* experiments were reported.

¹The General Environmental Technos Co., Ltd., Azuchimachi, Chuo-ku, Osaka, 541-0052, JAPAN (ishida hiroshi@kanso.co.jp)

²Research Institute of Innovative Technology for the Earth, Kizugawadai, Kizugawa, Kyoto, 619-0292, JAPAN

³Seto Marine Biological Lab., Field Science Education and Research Center, Kyoto University, Shirahama, Nishimuro, Wakayama, 649-2211, JAPAN

TREND OF OCEAN ACIDIFICATION IN THE WESTERN NORTH PACIFIC M26 <u>Ishii, Masao</u>^{1,2}, Takashi Midorikawa¹, Shu Saito¹, Takayuki Tokieda¹, Daisuke Sasano¹, Akira Nakadate^{1,2}, and Hisayuki Y. Inoue³

It is highly important to clarify the present status of ocean acidification by observations. Global Environment and Marine Department of JMA and MRI have been collaboratively continuing routine atmospheric/oceanic CO_2 measurements in the western North Pacific. These measurements in the tropical and subtropical zones along 137°E include pCO_2 in air and in seawater in January-February since early 1980s and in June-July since 1990, DIC section by coulometry since 1994, pCO_2 and sections of DIC and pH by spectrophotometry four times per year since 2003.

While large meridional gradients and seasonal variations in these parameters and temperature have been observed in surface layer of the northern subtropical zone ($\Delta p CO_2 sw = ca$. 90 μ atm, $\Delta NDIC$ (S=35) = ca. 35 μ mol kg⁻¹, ΔpH at SST = ca. 0.10, and $\Delta SST = ca$. 10 °C), the range of temporal and spatial variations in NTA calculated from these parameters is smaller (ca. 10 μ mol kg⁻¹). Increasing rates of $pCO_2 sw$ in January-February since 1983 and that in June-July since 1990 are both nearly equivalent to the increasing rate of atmospheric CO_2 during the same period. Such an increasing trend is also seen for NDIC in the water column above 26.8 σ_θ at around 30°N, where increasing rate of NDIC ranges from +1.0 to +1.5 μ mol kg⁻¹ yr⁻¹. These results strongly suggest that pH in the upper layer of the western North Pacific tropical and subtropical zones is decreasing at a mean rate of ca. -0.015 per decade.

¹Geochemical Research Department, Meteorological Research Institute, 1-1 Nagamine, Tsukuba, Ibaraki, 305-0052 Japan (mishii@mri-jma.go.jp)

²Global Environment and Marine Department, Japan Meteorological Agency, 1-3-4 Otemachi, Chiyoda-ku, Tokyo, 100-8122 Japan

³Graduate School of Environmental Science and Faculty of Environmental Earth Science, Hokkaido University, Kita-10, Nishi-5, Kita-ku, Sapporo, 060-0810 Japan

EFFECTS OF HIGH CO₂ ON DEEP-SEA FISHES T26

Ishimatsu, Atsushi, Masahiro Hayashi, and Yuki Kojima

This study aims to evaluate the effects of CO₂ plumes created by deep-sea CO₂ injection on deep-sea fishes. Three species of deep-sea fishes have been obtained alive from the depth of 380 m at a deep-water withdrawal facility in Toyama, Japan. They can be kept in laboratory conditions for several months under the atmospheric pressure at 1-2°C, and used for in vivo CO₂ exposure experiments under the atmospheric and elevated pressures. We have developed an experimental setup to expose deep-sea organisms to high CO₂ conditions under pressures of up to 20 MPa at 1-2°C, which also allows us to record electrocardiogram to monitor heart rate.

Under the atmospheric pressure, a deep-sea fish *Careproctus trachysoma* (Liparididae) showed 100% mortality within 48 h at 30,000 \square atm CO₂ conditions and 17% within 72 h at 20,000 \square atm. The other two species (*Allolepis hollandi* and *Malacocottus gibber*) showed mortalities similar to those we had

found for shallow-water fishes (mortality occurred mostly above 50,000 \square atm). The ongoing high-pressure experiment indicates the possibility that the elevated pressure of 6 MPa could reduce CO_2 tolerance of *C. trachysoma*. Video recordings demonstrated depressed gill ventilatory frequency in hypercapnic conditions under the atmospheric and 6 MPa conditions, in contrast to hyperventilatory responses to CO_2 known for most shallow-water fishes. Preliminary electrocardiogram recordings under 10 MPa demonstrated a rapid decline in response to 20,000 atm exposure (fish died at 13 h). Thus, some deep-sea fishes might be more sensitive to CO_2 than shallow-water species, particularly under high pressure conditions.

Institute for East China Sea Research, Nagasaki University, Nagasaki 851-2213, Japan (a-ishima@nagasaki-u.ac.jp)

IMPACTS OF OCEAN ACIDIFICATION ON BENTHIC PRIMARY PRODUCERS: LESSONS FROM A COLD ${\rm CO_2}$ VENT SITE ${\rm M27}$

Kerrison, Philip¹, David Suggett², Leanna Hepburn², and Michael Steinke²

Benthic autotrophs represent a small yet important fraction of total aquatic primary productivity. Whilst ocean acidification has been projected to impact the viability of calcifying organisms, the effect on photosynthesis and the production of the climate relevant trace gas dimethyl sulphide (DMS) is poorly understood. Previous studies speculated that high-CO2 conditions will (1) release autotrophs from the need to divert photosynthetic energy to carbon-concentrating mechanisms and (2) reduce the overall production of DMS and its precursor dimethylsulphoniopropionate (DMSP). Here we present data from a field study conducted on the island of Ischia in the Bay of Naples in May 2008. This site is characterised by cold volcanic vents of 90% CO₂ that create a pH gradient of about 6.0 to 8.2 along 300 m of shoreline. Recent evidence suggests that natural CO₂ vents can be used to reveal ecosystem effects of ocean acidification. We deployed two *in situ* pH loggers to describe the spatio-temporal variation in pH and conducted a survey of DMSP and DMS production in seaweeds and benthic microalgae along the pH gradient. A characterisation of primary productivity and photosynthetic efficiency complements our data set. Our results address changes in marine benthic communities in a future ocean scenario and their possible effect on primary productivity and air-sea emissions of DMS.

¹University of Essex, Department of Biological Sciences, Colchester, CO2 8JP, U.K. (pkerri@essex.ac.uk).

EFFECTS OF HIGH CO₂ CONCENTRATIONS ON THE PHAEODACTYLUM TRICORNUTUM (BACILLARIOPHYCEAE) AND NODULARIA SPUMIGENA (CYANOPHYTA) GROWTH AND ORGANIC MATTER PRODUCTION T27 Kosakowska, Alicja, and Janusz Pempkowiak

The coastal seas and estuaries are especially vulnerable to effects of increased CO_2 in the air. The extend by which production of organic matter is enhanced under increased CO_2 availability is of utmost importance for the areas. Also effects of chemical changes in the river run-off are important and require clarification.

In order to investigate the influence of increased CO₂ concentration in the atmosphere on the phytoplankton growth and organic matter production at three CO₂ concentrations, mimicking the present day (360ppm), the near future (600ppm), and a would be (1000ppm) conditions a series of experiments were performed. The influence was measured using two phytoplankton species, namely diatom *P.tricornutum* and Baltic cyanobacterium *N.spumigena*.

The growth rate and chlorophyll content were enhanced when *P.tricornutum* was grown at high CO₂ with respect to normal CO₂. The growth at 600 and 1000ppm CO₂ was 1,5 and 2 times higher as compare to 360ppm CO₂. On the other hands the production of chlorophyll and cells growth of *N.spumigena* were both negatively affected at 600 and 1000 CO₂ ppm during a 5 days long

²University of Essex, Department of Biological Sciences, Colchester, CO2 8JP, U.K.

incubations, and positively affected at 600ppm CO₂ after 7 days cultivation. Significant differences in chlorophyll content were observed between cells exposed to 600ppm CO₂ enriched air and the control, as were in the DOC concentrations in sea water used as culture medium.

The contrasting influence of the high CO₂ on the growth of these species suggests that CO₂ may alter algal community composition and species succession under actual high CO₂ stress.

Institute of Oceanology, Polish Academy of Sciences, Powstańców Warszawy 55, Sopot, Poland (akosak@jopan.gda.pl)

COCCOLITHOPHORE COMMUNITY RESPONSE TO OCEAN ACIDIFICATION M28 Krug, Sebastian A., Birte Matthiessen, Ulf Riebesell, and Ulrich Sommer

The effects of ocean acidification on the calcification and photosynthesis of coccolithophores has been a focus of research for more than a decade. So far most studies have relied on experiments with single species strains, indicating taxon-specific differences in the responses to rising atmospheric pCO₂. While environmental perturbations can have profound impacts on the diversity of an ecosystem and it has been hypothesised that diverse systems are able to compensate perturbations more efficiently than species poor systems. Hence, this experiment focuses on the capacity of a multispecies assemblage to absorb stress and re-organize while undergoing change. This has been addressed by exposing an artificial community consisting of *Emiliania huxleyi*, *Gephyrocapsa oceanica* and *Coccolithus pelagicus braarudii* to elevated CO₂ and comparing its response to that of the individual species in monocultures. Preliminary results of fully factorial tested effects of decreasing pH/increasing pCO₂ show significant treatment differences in the biomass production of monocultures compared to the coccolithophore assemblage. In terms of biomass production the community was able to compensate the CO₂/pH stress and underwent a shift from a *Gephyrocapsa oceanica* dominated assemblage at low pCO₂ towards an *Emiliania huxleyi* dominated system at a high pCO₂.

Leibniz Institute of Marine Sciences, IFM-GEOMAR, Duesternbrooker Weg 20, 24105 Kiel, Germany. (skrug@ifm-geomar.de)

REPRODUCTIVE PHYSIOLOGY OF SEA URCHIN IN A HIGH-CO₂ WORLD Kurihara, Haruko Rui Yin and Atsushi Ishimatsu

Calcifying marine organisms are the most vulnerable to reductions of calcium carbonate (CaCO₃) saturation state. Although previous studies revealed effects of high CO₂ seawater on calcification rate, growth rate and physiology of adult calcifiers, little is known about impacts on early development and reproduction, which is crucial to understand CO₂ effects in a population level, since these are usually the most sensitive stages of a life cycle and directly affect population size. Additionally, it is critical to understand their synergetic effects of high CO₂ and temperature on marine organisms.

We studied the reproductive physiology of sea urchin $Hemicentrotus\ pulcherrimus$ reared under high CO_2 (1,000 ppm CO_2) and high CO_2 + high temperature (1,000 ppm CO_2 + 2° C) condition for 8 months. We found that both conditions impacted the physiology, GSI (gonad wet weight / wet weight) and gonad development of the sea urchin, although these conditions did not affect the survival and growth rate (size) of sea urchin. These results suggest that a future high- CO_2 world will potentially affect the reproductive success of sea urchin. Together with our previous findings of negative impacts of increased seawater CO_2 on the early development, we suggest that sea urchin populations will be threatened in a high- CO_2 world.

Institute for East China Sea Research, Nagasaki University, Nagasaki, Japan (harukoku@e-mail.jp)

EFFECTS OF INCREASED CO2 AND TEMPERATURE ON TRACE-ELEMENT BIOACCUMULATION IN EGGS OF THE COMMON CUTTLEFISH, SEPIA OFFICINALIS M29

<u>Lacoue-Labarthe</u>, Thomas^{1,2}, Sophie Martin¹, François Oberhänsli¹, Jean-Louis Teyssié¹, Ross Jeffree¹, and Paco Bustamante²

Cephalopods play a key role in many marine trophic networks and constitute alternative fisheries resources, especially given the ongoing decline in finfish stocks. Cephalopods die after mating, which means that their population dynamics are highly dependent on the hatching success of their eggs. Along the European coast, the eggs of the cuttlefish Sepia officinalis are characterized by an increasing permeability of the eggshell during development, which leads to selective accumulation of essential and non-essential elements in the embryo. Temperature and pH are two critical factors that affect the metabolism of marine organisms, but very few studies have considered the synergistic effect of both factors on trace-element bioaccumulation. In this study, we are testing the effects of pH and temperature through a crossed (3x2) laboratory experiment. Cuttlefish eggs are being reared at normal pH (8.10) and at two lower pH's (7.85 and 7.60); all three pH conditions are being studied at two temperatures (16°C and 19°C). Eggs are being exposed to ⁴⁵Ca, ⁵⁴Mn, ⁶⁰Co, ⁶⁵Zn, ¹⁰⁹Cd, ^{110m}Ag, ¹³⁴Cs, and ²⁴¹Am tracers that were dissolved in seawater in order to assess their uptake kinetics and spatial redistribution within the egg compartments and how that varies during the embryonic development time. Temperature and pH could affect embryo metabolic rates and chemical properties of eggshell components, which could lead to shifts in a) the accumulation of essential elements (Ca, Co, Mn, Zn) that are needed by the embryo and b) the capacity of the eggshell to shield against penetration of nonessential or toxic elements (Ag, Cd, ¹³⁴Cs, ²⁴¹Am).

EFFECT OF HIGH ${\rm CO_2}$ ON THE PHYSIOLOGY AND GROWTH OF THE HERMATYPIC CORAL SERIATOPORA HYSTRIX IN NANO-SIZED CHEMOSTAT T29

Leblud, J., A. Batigny and Ph. Grosjean

Growth of hermatypic coral *Seriatopora hystrix* was studied in a new original artificial chemostat. We succeed to maintain and growth small nubbins of 0.23±0.05g of the hermatypic coral *S. hystrix* in nano-sized chemostats (called "nanocultures") of 400 mL each during 3 weeks. Main physicochemical parameters are monitored and controlled using microcomputers. This includes temperature, salinity, pH, alkalinity, omega aragonite and calcium concentration. Artificial light of 250±30 μE.m⁻².s⁻¹ on a 12h/12h light/dark cycle and a temperature of 27.0±0.1°C, salinity of 34±1, pH of 8.2±0.2, alkalinity of 2.9±0.3meq.kg⁻¹ and [Ca⁺⁺] of 410±10 mg.kg⁻¹ give a growth rate of 1.1±0.5% per day for the nubbins, i.e., an increase in weight of more than 25% during the experiment.

The study of variation of the physico-chemical parameters, including true duplicates or triplicates, are possible with our system. In the near future, we propose to investigate the effect of increasing CO₂ on growth and physiology of *S. hystrix*, his symbiotic zooxanthellae and his epibiontic bacterial colonies using these nanocultures. Relatively axenic conditions, measurement of the complete balance of inorganic/organic carbon in each chemostat, and concentration of the organic molecules released by the coral due to the extremely small volume potentially offer new insights on coral growth and physiology.

Académie Universitaire Wallonie-Bruxelles, Université de Mons-Hainaut, Laboratoire d'Ecologie Numérique des Milieux Aquatiques, 6 Avenue du Champ de Mars, 7000 Mons, Belgium. (julien.leblud@umh.ac.be)

¹ Marine Environmental Laboratories, International Atomic Energy Agency, Monaco.

² Littoral Environnement et Sociétés, CNRS-Université de la Rochelle, La Rochelle, France.

REGULATION OF NITROGEN METABOLISM IN THE MARINE DIAZOTROPH TRICHODESMIUM IMS101 UNDER VARYING TEMPERATURES AND ATMOSPHERIC CO₂ CONCENTRATIONS. M30

<u>Levitan, Orly</u>¹, M. Christopher Brown², Stefanie Sudhaus³, Douglas Campbell², Julie LaRoche³, and Ilana Berman-Frank¹

Trichodesmium spp., the dominant cyanobacterial nitrogen-fixer in the tropical and sub-tropical oligotrophic oceans, responds to elevated pCO₂ (from current to 900 □atm) with a 3-4 fold increase in nitrogen fixation rates. Here we examined regulation of N₂ fixation at the levels of transcript (nifH mRNA copies), protein, and activity (acetylene reduction) over ranges in both CO₂ and temperature [250 (low), 400 (ambient) and 900 (high) \square atm CO₂ at 31 $^{\circ}$ C and 25 $^{\circ}$ C]. While the amount of nitrogenase protein remained comparable across all temperature and CO₂ combinations, both nifH transcription and activity levels were significantly influenced by CO2 concentrations. nifH transcription in cultures acclimated to high CO₂ (at both temperatures) declined throughout the photoperiod, contrasting with the characteristic diel pattern of nitrogen-fixation which was markedly higher under elevated CO₂. Estimated catalytic rates for both nitrogenase and the subsequent nitrogenassimilatory enzyme, glutamine synthetase, revealed a parallel diel pattern. At high CO₂ these enzymes operated at overall catalytic rates of over 60% of the *in-vitro* k_{cat} values while under ambient and low CO₂ these enzymes operated at only ~30% of k_{cat}. At elevated CO₂, the lower transcript levels combined with faster enzymatic catalytic rates may indicate enhanced stability and lower turnover of the protein. Thus, acclimation of Trichodesmium to elevated CO₂ lowers the energy costs for maintenance of the nitrogenase protein pool, while reallocation of energy and resources from competing pathways such as the Carbon Concentrating Mechanism and the Mehler reaction can stimulate the higher N₂ fixation and ultimately growth rates.

CARBON CYCLE PERTURBATIONS DUE TO MODIFIED CARBON FLUX AND FUTURE CO $_2$ UPTAKE IN THE NORTHERN INDIAN OCEAN T30

Mahanta, Chandan

Large uncertainty exists concerning uptake of CO₂ by Indian Ocean due to insufficient knowledge of processes controlling carbonate chemistry. Settling of particulate organic carbon decreases total C and pCO₂ of surface layers, whereas removal of carbonate from surface waters aids in increasing atmospheric CO₂ by shifting carbonate equilibrium. The method in calculating sea-air CO₂ exchange was based on flux equation considering difference between partial pressure of atmosphere and sea surface that determine the direction of CO₂ exchange. While gas transfer velocity depends on wind velocity on sea surface, monsoonal episodic pulse of about 5% global particulate riverine C load received by Bay of Bengal shelf is important natural and human induced interactive forcing influencing future capacity of CO₂ assimilation. Bay of Bengal, a high productive area of the world oceans, has recorded high organic carbon fluxes of more than 3 g m⁻²y⁻¹ in sediment trap experiments. Air-sea CO₂ flux variability impact due to variations in entrained terrestrial input could be more pronounced than surface temperature or wind speed. Result showed, in the past, release was around 2 moles CO₂ m⁻² year⁻¹. Capacity of oceanic uptake continued to increase, and in recent times, the region absorbed up to 5 moles CO₂ m⁻² year⁻¹. The prediction result based on the A2 scenario in the year of 2050 showed that BOB waters can optimally uptake up to 15 moles CO₂ m⁻² year⁻¹, but would begin to drop due to decreasing of atmospheric CO₂ concentration until 2100. The B2 scenario showed a lower trend.

¹The Mina and Everard Goodman Faculty of Life Sciences, Bar Ilan University, Ramat Gan, 52900, Israel (levitao@mail.biu.ac.il).

²Department of Biology, Mount Allison University, Sackville, NB E4L 1G7, Canada.

³Department of Marine Biogeochemistry, IFM-GEOMAR, Leibniz Institute of Marine Sciences at Kiel University, Duesternbrooker Weg 20, 24105 Kiel, Germany.

Civil Engineering Department, IIT Guwahati, Guwahati-781039, Assam, India. Phone: 91-361-2691000, Fax: 91-361-2582440, Email: mahanta iit@yahoo.com

DRASTIC DECREASE IN COLD WATER CORAL CALCIFICATION AS RESPONSE TO LOWER PH M31

Maier, Cornelia^{1,2}, Jan Hegeman¹ and Markus G. Weinbauer²

The cold-water coral *Lophelia pertusa* is one of the few species able to build reef-like structures and a 3-dimensional coral framework in the deep oceans. Such deep coral bioherms are likely among the first to be affected by ocean acidification i.e. decrease in pH due to an increase in pCO2. During a cruise in the Skagerrak, calcification rates and response to lowering pH has been studied directly onboard using cold-water corals that have been freshly collected with a video-equipped box corer. Calcification rates were assessed using 45 CaCl₂. Experiments were conducted at ambient pH and pH lowered by 0.15 and 0.3 units. Calcification rates were higher than expected with an average of 0.05 % $^{4-1}$ ±0.01 S.E. (normalised to initial skeletal weight) for bulk calcification of small branches. Highest calcification rates were found in youngest polyps, which had calcification rates of up to 1 % $^{4-1}$ and average calcification rates of 0.11 % $^{4-1}$ ± 0.02 S.E. (extremes removed). Lowering the pH by 0.15 and 0.3 pH units reduced coral calcification by 30 % and 56 %, respectively. Also, the effect of changes in pH (0.3 pH units lower than in ambient water) on calcification rate was stronger for fast growing, young polyps (59 % reduction) than for older polyps (40 % reduction). This first study on calcification rate and pH effects for *L. pertusa* implies that the young and fast calcifying corallites will be influenced most negatively by ocean acidification.

EFFECTS OF OCEAN ACIDIFICATION ON EARLY LIFE STAGES OF SEA BREAM AND SEA BASS T31

<u>Martin, Sophie</u>¹, Thomas Lacoue-Labarthe², Fanny Houlbreque¹, François Oberhänsli¹, Jean-Louis Teyssié¹, Steeve Comeau³, Florence Boisson¹, Jean-Pierre Gattuso³, James Orr¹ and Ross Jeffree¹

The gilthead sea bream, *Sparus aurata*, and the European sea bass, *Dicentrarchus labrax*, provide the largest economic sector for fish aquaculture along the Mediterranean and Eastern Atlantic coasts, with global production of 108,000 tons of sea bream (\$595 million US) and 60,000 tons of sea bass (\$386 million) in 2006. Despite their commercial interest and their wide geographic distribution, we do not understand how ocean acidification will affect these species and in particular their early developmental stages that could be particularly vulnerable to elevated CO₂ partial pressure (pCO₂). We investigated the consequences of ocean acidification on the embryonic and larval stages of *S. aurata* and *D. labrax*. Fish eggs and larvae were maintained in Mediterranean seawater at normal surface conditions (pH=8.1, pCO₂=380 ppm) and at three low-pH, high CO₂ conditions: pH=7.9 (650 ppm), pH=7.7 (1100 ppm), and pH=7.5 (1800 ppm). Eggs and larvae were exposed to ⁴⁵Ca, ⁵⁴Mn, ⁶⁰Co, ⁷⁵Se, ^{110m}Ag, ¹³⁴Cs, ²⁴¹Am, ⁶⁵Zn, and ¹⁰⁹Cd dissolved in seawater in order to assess the uptake kinetics of these elements from the spawning date to the end of the first week of larval development. Preliminary results reveal that ocean acidification may alter incorporation of trace elements in eggs and larvae. Both essential metals (Co and Zn) and non-essential metals (Ag and Cd) were taken up less efficiently by the eggs at reduced pH relative to normal pH, suggesting potential effects on fish physiological functions and on retention of toxic elements.

¹ Royal Netherlands Institute for Sea Research (NIOZ), Dept of Biol. Oceanography, BP 59, 1790 AB Den Burg, the Netherlands

²Laboratoire d'Océanographie de Villefranche (LOV), Microbial Ecology and Biogeochemistry Group, CNRS-UPMC, UMR 7093, BP 28, 06234 Villefranche-sur-mer, France, (maier@obs-vlfr.fr)

¹ Marine Environmental Laboratories, International Atomic Energy Agency, Monaco.

² Littoral Environnement et Sociétés, CNRS-Université de la Rochelle, La Rochelle, France.

³ Laboratoire d'Océanographie de Villefranche, CNRS-Université de Paris VI, Villefranche-sur-Mer, France.

HIGH CO₂ RETAINED FRACTION IN OCEAN SEQUESTRATION ESTIMATED BY A HIGH RESOLUTION MODEL M32

Masuda, Yoshio¹ and Yasuhiro Yamanaka^{1,2}

In the CO₂ ocean sequestration, CO₂ retained fraction is one of the most important topics. It shows how long CO₂ injected into the ocean is isolated from the atmosphere. The retained fraction of the injected CO₂ calculated in previous studies are around 70% in average after 200 years (IPCC report) which are calculated by relatively coarse resolution models with the horizontal resolution larger than one degree. We considered that the retained fraction should be small, because 200 years are much shorter than the time scale (~2000 years) of deep ocean circulation. We estimated it using a high resolution model with the horizontal resolution of 0.1 degree. The fine grid mesh explicitly represent effects of mesoscale eddies on CO₂ transport and dilution. The injected CO₂ is kept in smaller ocean volume than that obtained from coarse resolution models. Only 0.4% of the injected CO₂ is released into the atmosphere after 200 years, although about half of the injected CO₂ is horizontally transported to the outside of the narrow model domain. We supposed that the difference in the retained fraction between high and coarse resolution models is caused by difference in horizontal CO₂ transport. Difference in simulated western boundary current regions also may effect on the difference.

¹Hokkaido University, N10, W5, Kita-ku, Sapporo, Hokkaido 060-0810, Japan (y-masuda@ees.hokudai.ac.jp)

ENHANCED BIOLOGICAL CARBON CONSUMPTION IN A HIGH CO₂ OCEAN: A REVISED ESTIMATE OF THE EFFICIENCY OF THE OCEANIC CARBON UPTAKE T32 Matear, Richard¹ and Ben McNeil²

Using recent empirical measurements of enhanced biological carbon consumption in a high CO₂ ocean (Riebesell et al., 2007), we have investigated the potential for this effect to increase the carbon uptake by the oceans in an ocean general circulation model (OGCM) and a data-calibrated two box model. Our results suggest an increased oceanic carbon uptake of 46 GtC up to the year 2100, which is less than half the value estimated by Riebesell et al., (2007). The reduced oceanic CO₂ uptake comes about through a more realistic parametrization of the re-supply of carbon from the interior ocean back to the surface than originally estimated from Riebesell et al., (2007) simple model. The OGCM simulations further reveals that the oceanic circulation modifies the distribution of carbon whereby some regions like the eastern equatorial Pacific and Southern Ocean experience reduced carbon uptake despite a local increase in biological carbon export. The pooling of the exported carbon in these regions allows the re-supply of carbon rich sub-surface waters to exceed the enhanced biological carbon export of the high CO₂ world.

INCREASED CO2 UPTAKE IN THE ARCTIC OCEAN: HOW SEA ICE LOSS WILL IMPACT OCEAN ACIDIFICATION M33

Mathis, Jeremy T.1 and Nicholas R. Bates2

The Chukchi shelf is the site of some of the highest rates of primary production in the oceans. Rates of net community production over the shelf can be as high as 2,000 mg C m-2 d-1, and averaged 800 mg C m-2 d-1 across the entire shelf. Using conservative tracers to construct a carbon mass balance,

²CREST, JST, Sanbancho 5, Tokyo, 102-0075, Japan.

¹Centre for Australian Weather and Climate Research (CAWCR)

²University of New South Wales

we found that 10% of the DIC consumed during net community production was converted to DOC and 15% was converted to suspended POC. The remaining 75% was exported from the mixed layer as sinking organic particles. At the termination of the bloom, most of the organic matter had been exported from the mixed layer, leaving surface waters undersaturated with respect to atmospheric CO2. Before reequilibration can occur with the atmosphere the surface waters are transported offshore, under the ice cap which further inhibits air-sea exchange and maintains low pCO2 levels in the surface waters of the deep Arctic Basin. However, in recent years, the extent of summer sea ice has retreated further into the Arctic Ocean with a 30% reduction observed in summer of 2007. This "opening" of the central Arctic has lead to an increased uptake of CO2 into the central basin, thus lowering the pH of surface waters in the region. Here, we show the distributions of pH over the Chukchi Sea and adjacent Canada Basin in 2002 and 2004 and discuss the impacts that future sea ice loss will have on ocean acidification in the Arctic.

CO₂ VS. HCL FOR THE ADJUSTMENT OF SEAWATER pH T33

McGraw, Christina M.¹, Catriona L. Hurd², Christopher D. Hepburn², Kim I. Currie³, John A. Raven⁴, and Keith A. Hunter¹

The impact of acidification on the world's oceans is often predicted through the culture of marine organisms under controlled conditions. Physiological responses are monitored as seawater pH is adjusted by additions of HCl or bubbling of CO₂ gas. Both methods easily achieve target pH levels, but calculation of CO₂ equilibrium speciation shows that they have significantly different impacts on seawater chemistry and the concentration of dissolved inorganic carbon species. When CO₂ is used to lower the seawater pH, the total alkalinity (A_T) remains constant while the total inorganic carbon (C_T) increases. When HCl is used to lower the seawater pH, C_T remains constant while A_T decreases. Simulations show that the resulting differences in [CO₂(aq)], [CO₃²⁻], and *p*CO₂ for these two methods of acidification are small. However, [HCO₃-] is significantly higher for seawater adjusted with CO₂ compared to seawater adjusted with HCl. This difference in CO₂ speciation between the two methods of acidification may be significant for organisms that use HCO₃- for photosynthesis. These results indicate that pH and CO₂ speciation must both be considered when designing culture experiments. In addition, the calculations show that seawater adjusted with CO₂ more closely mimics the changes in seawater carbon chemistry that are expected with rising atmospheric CO₂ levels.

EARLIER SOUTHERN OCEAN ACIDIFICATION FROM SEASONAL AMPLIFICATION OF ANTHROPOGENIC CO₂ UPTAKE M34

McNeil, Ben I.¹ and Richard J. Matear²

Southern Ocean acidification via anthropogenic CO₂ uptake is expected to be detrimental to multiple calcifying plankton species by lowering the concentration of carbonate ion (CO₃²⁻) to levels where calcium carbonate (both aragonite and calcite) shells begin to dissolve. Although seasonal variability has been suggested to hasten the onset of aragonite under-saturation (Orr et al., 2005), observational

¹ University of Alaska Fairbanks, School of Fisheries and Ocean Sciences, 245 O'Neil BLDG Fairbanks, AK, USA, 99775 (jmathis@sfos.uaf.edu).

² Bermuda Institute of Ocean Sciences, St. George's, GE 01, Bermuda, (nick.bates@bios.edu).

¹Department of Chemistry, University of Otago, PO Box 56, Dunedin, New Zealand (cmcgraw@chemistry.otago.ac.nz)

²Department of Botany, University of Otago, PO Box 56, Dunedin, New Zealand

³National Institute for Water and Atmospheric Research Ltd, Centre of Excellence for Chemical and Physical Oceanography, Department of Chemistry, University of Otago, PO Box 56, Dunedin, New Zealand.

⁴Division of Plant Sciences, University of Dundee at SCRI, Scottish Crop Research Institute, Invergowrie, Dundee DD2 5DA, U.K.

evidence in the Southern Ocean has been lacking. We present a large-scale Southern Ocean observational analysis that examines the seasonal magnitude and variability of CO_3^{2-} and pH. Our analysis shows an intense winter-time minimum in CO_3^{2-} south of the Antarctic Polar Front and when combined with projected anthropogenic CO_2 uptake will induce aragonite under-saturation as early as the year 2030 and no later than 2038. Some prominent calcifying plankton, in particular the Pteropod species Limacina helicina, have important veliger larval development during winter and will have to experience detrimental carbonate conditions much earlier than previously thought, with possible deleterious flow-on impacts for the wider Southern Ocean marine ecosystem. Our results highlight the critical importance of understanding seasonal carbon dynamics within all important calcifying marine ecosystems such as continental shelves and coral reefs, since it may potentially hasten the onset of future ocean acidification.

¹Climate Change Research Centre, Faculty of Science, University of New South Wales, Sydney, NSW, Australia.

A partnership between CSIRO and the Bureau of Meteorology, Hobart, TAS, Australia

EVOLUTION OF PRE-ADAPTATION TO FUTURE OCEAN ACIDIFICATION: IDENTIFYING PHYSIOLOGICAL CHARACTERISTICS OF SURVIVORS T34

Melzner, Frank ¹, Magdalena A. Gutowska², and Martina Langenbuch ¹

Several recent reviews have summarized potential dangers of future ocean acidification to marine animal survival and performance. However, we are still far from understanding the genetic and physiological mechanisms that provide certain groups of high-power animals (fish, cephalopods, some crustaceans) with a relative degree of tolerance towards simulated ocean acidification. Employing an evolutionary approach, this paper will synthesize information on how the development of high-power ecotypes resulted in the co-evolution of a relative acidification tolerance. High-power organisms experience high pCO₂s in their body fluids (1,000-8,000 ppm) under control conditions and during exercise already in today's ocean. Thus, contrary e.g. to unicellular ocean organisms, who are bathed in an extracellular fluid (the ocean) that has a pCO₂ of <400 ppm, high-power animals have already had a long time to evolve physiological mechanisms to live with high internal pCO₂s. Using the physiological literature of the last 40 years in combination with recent investigations on acid-base regulation in marine invertebrate animals in our laboratories, we will classify species groups (teleost fish, cephalopods, crustaceans, bivalves, echinoderms, cnidarians) according to key physiological parameters (ion exchange capacities and mechanisms, metabolic rate fluctuations, CO₂ exchange rates and body [CO₂] variations), to elaborate common patterns. We also will address the pressing question, whether the currently anticipated speeds of future acidification will leave enough time for evolution of such traits that promote tolerance to high ocean CO₂.

ARE EASTERN BOUNDARY UPWELLING SYSTEMS GOOD ANALOGUES FOR A FUTURE HIGH CO2 OCEAN? M35

Monteiro, P.M.S.¹ and R.G.J. Bellerby²

Eastern boundary upwelling systems are characterized by waters with elevated pCO2 (500 – 1500ppm) and low pH (ranges?) which span the envisaged carbonate system characteristics resulting from future ocean acidification. This is especially the case where the shelf edge boundary conditions for the upwelled waters are derived from Equatorial OMZ thermocline waters. Advection of these waters over organic rich shelf sediments where aerobic and anaerobic respiration fluxes of CO2 prior to upwelling, create low pH (pH < 7.6) conditions in surface and sub-thermocline waters. Observational data from the Benguela upwelling system show direct effects of these conditions

²Centre for Australian Weather and Climate Research (CAWCR)

¹IFM-GEOMAR, Leibniz Institute for Marine Sciences, Kiel, Germany (fmelzner@ifm-geomar.de)

² Alfred-Wegener-Institute for Polar and Marine Research Bremerhaven, Germany

through malformations of coccolithophorid shells in the surface layer (Giraudeau et al., 1993). The impacts of elevated CO2 may extend to other parts of the carbon and nitrogen biogeochemistry and possibly early life stages of calcifying and non calcifying organisms.

We have re-analyzed historical and more recent data with a view to establishing the extent and characteristics of variability of this natural acidification in the Benguela upwelling system.

¹CSIR, PO Box 320, Stellenbosch, South Africa, pmonteir@csir.co.za

IMPACTS OF OCEAN ACIDIFICATION ON SCLERACTINIAN CORAL SKELETAL INTEGRITY T35

Mooney, Ann¹, David Kline², and Kenneth Anthony²

Carbon dioxide emissions are well-documented to have risen exponentially since pre-industrial times. The rise of emissions has numerous effects on our environment, and coral reefs are emerging as one of the most susceptible ecosystems. Increasing emissions are resulting in warmer and more acidified seas that may compromise coral calcification and growth. Among many ecological and physiological effects, the ways in which the skeletal integrity of reef-building corals may be compromised by the rise of CO_2 must be addressed. The following study examines how the strength of the coral skeleton is affected by rising atmospheric CO_2 . Samples from highly abundant corals representing differing skeletal characteristics were exposed to varying levels of CO_2 to represent future reef scenarios. Those corals exposed to high CO_2 levels grew similar amounts as the control corals; however, they broke with much less energy. This study sought to provide managers and policy-makers with quantifiable data regarding skeletal strength.

¹National Oceanic and Atmospheric Administration, Papahanaumokuakea Marine National Monument. 6600 Kalaniana'ole Hwy suite 300, Honolulu, HI 96825 USA (ann.mooney@noaa.gov) ²Centre for Marine Studies, University of Queensland, St. Lucia, QLD 4072 Australia

LONG TERM NEUTRALIZATION OF ANTHROPOGENIC CO₂: THE ROLE OF CHANGING BASIN-TO-SHELF PARTITIONING OF CARBONATE BURIAL M36 Munhoven, Guy

The longterm implications of the ocean uptake of CO₂ released into the atmosphere by human activity on the partitioning of the global carbonate burial between shelf and pelagic sea-floor sediments is analyzed with MBM, a multi-box model of the carbon cycle in the ocean and atmosphere, fully coupled to the transient early diagenesis model MEDUSA. MBM-MEDUSA includes separate schemes for the accumulation of carbonate on the shelf (coral-reefs and banks) and in the deep-sea sediments; carbonate accumulation by coral-reefs is dependent on the saturation state of aragonite in the surface ocean. An emission scenario following a logistic function is adopted to prescribe the release of CO₂ from fossil fuel burning over the next centuries until depletion of resources (assumed to be 4240 GtC). Under that scenario, the net carbonate accumulation by coral-reefs rapidly decreases during the 21st century. Accumulation rates will be reduced by more than 90% in 2100 and almost completely vanish by the year 2200. Accumulation in coral-reef environments then slowly resumes, but even 20,000 years later, it only reaches 80% of the Holocene level. A complete feedback analysis regarding the role of the resulting change in the partitioning of carbonate burial between the shelf and deep-sea sediments for atmospheric CO₂ concentrations will be presented.

Laboratoire de Physique Atmosphérique et Planétaire, Université de Liège, B-4000 Liège, Belgium (Guy.Munhoven@ulg.ac.be).

²BCCR, 55 Allegaten, Bergen, Norway

AN EXPERIMENTAL STUDY SITE FOR OCEAN ACIDIFICATION AT THE UNIVERSITY OF WASHINGTON, FRIDAY HARBOR LABORATORIES. T36

Murray, James W. and Terrie Klinger²

We are leading an effort at the University of Washington to design and construct an experimental study facility for ocean acidification at the Friday Harbor Laboratories (FHL). This facility will allow experimental studies of the response of marine organisms to changes in ocean pH and carbonate saturation in an environmentally-relevant setting. We anticipate that this facility will grow to become a site accessible for US and Canadian scientists to conduct experimentation on the biological and ecological impacts of ocean acidification. FHL is uniquely suited for this facility because it is an established center for research in marine biology, ecology and oceanography with a stellar international reputation. Its location allows experimentation on organisms native to the northeastern Pacific, which is predicted to show early, strong signs of acidification. This exciting new project brings together chemical and biological oceanographers from the Schools of Oceanography, Aquatic and Fisheries Sciences, Marine Affairs and The Friday Harbor Laboratories at the University of Washington. The initial centerpiece will be development of an experimental study site at the FHL that will incorporate both small scale lab based and larger scale mesocosm type experiments that will be used for studies of both ocean acidification and ocean warming. A planning workshop will be held at Friday Harbor Laboratories on 28, 29 August and the recommendations will be presented at this Symposium.

MODELLING THE UPTAKE AND RELEASE OF CARBON DIOXIDE IN THE BALTIC SEA SURFACE WATER M37

Omstedt, A, Erik Gustafsson, and Karin Wesslander

The present work is the first attempts within GEWEX/BALTEX phase II to model the uptake and release of carbon dioxide in the Baltic Sea. The mathematical formulation is kept as simple as possible, but several aspects of the dynamics are included. The model system captures major physical-chemical and biological response patters as evaluated from observations from the central Baltic Sea and the modeling illustrates the need for fractional nutrient release in the photic zone for consistency with the CO₂ observations.

The carbon dioxide exchange between Baltic Sea and atmosphere are besides from carbon chemistry, strongly dependent on biologic production and mineralization. Modeling of long-term variations in partial pressure of carbon dioxide indicates stable conditions before industrialization and net release to the atmosphere. During modern industrialization era with eutrophication the modeling illustrates that the seasonal variability increased and the net release to the atmosphere decreased. The study illustrates that observed and modeled partial pressure of carbon dioxide gives important knowledge about time evolution and size of primary production. The modeling illustrates also the need for fully coupled physical-chemical-biological ecosystem models where the carbon chemistry is included. Improved knowledge about river and Skagerrak input of organic and inorganic carbon as well as alkalinity are needed.

Department of Earth Sciences; Oceanography, University of Gothenburg, Sweden

¹ School of Oceanography, Box 355351E, University of Washington, Seattle WA 98195-5351, USA (jmurray@u.washington.edu)

² School of Marine Affairs, University of Washington, Seattle WA 98105-6715, USA (tklinger@u.washington.edu)

VOLATILE ORGANIC COMPOUNDS (VOCs) IN SEA WATER UNDER ELEVATED CO₂ LEVELS DURING AN OFF-SHORE MESOCOSM EXPERIMENT T37

Orlikowska, Anna and Detlef Schulz-Bull

Volatile organic compounds (VOCs) are known to play an important role in the global climate and are involved in a number of photochemically induced atmospheric reactions. The processes responsible for formation and degradation of VOCs are complex and are affected by various factors among them modifications in seawater chemistry.

In the current study water samples are analysed in order to investigate the influence of various parameters - especially elevated CO_2 levels, changes in irradiance and temperature - on concentration and composition of volatile hydrocarbons. Moreover stable carbon isotope analysis is performed to study $\delta 13C$ values. While VOCs are ubiquitous trace gases released into the environment from natural and anthropogenic sources this technique can be used to make a distinction between their various origins as well as serving as valuable tool to distinguish biodegradation from physical, nondegradative processes. The seawater samples were taken during the SOPRAN off-shore mesocosm experiment in the central Baltic Sea in July 2007 and 2008. In an attempt to investigate CO_2 effects on marine ecosystem acid treatment is used to induce different p CO_2 environments in the mesocosms. The analysis is carried out by a purge and trap system coupled to a gas chromatograph interfaced with mass spectrometer and isotope ratio mass spectrometer. Twenty five low molecular weight volatile halogenated organic compounds (VHOCs) along with dimethylsulfide (DMS) and isoprene as well as monocyclic aromatic compounds are determined. Relations between the compounds as well as the influence of physicochemical factors are presented.

Department of Marine Chemistry, Leibniz Institute for Baltic Sea Research Warnemünde (IOW), Seestraße 15, 18119 Rostock, Germany (anna.orlikowska@io-warnemuende.de)

ARCTIC ACIDIFICATION M38

Orr, James C.,¹, Sara Jutterström², Laurent Bopp³, Leif G. Anderson², Victoria J. Fabry⁴, Thomas Frölicher⁵, Peter Jones⁶, Fortunat Joos⁵, Ernst Maier-Reimer⁷, Joachim Segschneider⁷, Marco Steinacher⁵ and Didier Swingedouw⁸

Climate change in the Arctic will be amplified, leading to reduced sea-ice cover, warming and freshening of surface waters, and changes in vertical stratification. The Arctic Ocean will also undergo acidification. Previous modeling studies suggest that the coldest surface waters of the Southern Ocean will be the first to become undersaturated with respect to aragonite, the metastable form of calcium carbonate (CaCO₃), i.e., within 50 years under the IS92a scenario. However, those studies did not discuss the potential for similarly dramatic changes in the Arctic Ocean, owing to lack of gridded baseline data in that region. To assess future CaCO₃ saturation in the Arctic Ocean, we used recent data along two trans-Arctic sections as a baseline, to which we added projected 21stcentury perturbations in DIC, alkalinity, temperature, salinity, and nutrients from three coupled carbon-climate models forced under the IPCC SRES A2 scenario. In our projections, some Arctic Ocean surface waters become undersaturated with respect to aragonite within 10 years. On average, surface waters succumb to these conditions by 2050. By 2060, some surface waters become undersaturated with respect to calcite, the stable form of CaCO₃. At risk are pelagic and benthic marine calcifiers, including bivalve molluscs, a prominent species of the Arctic-shelf benthic community and a major food source for walruses, grey whales, and spectacled eiders. Owing to amplified Arctic climate change, which exacerbates effects from elevated CO2, undersaturated conditions detrimental to ecosystems will develop first in the Arctic Ocean, 10 to 30 years sooner than in the Southern Ocean.

¹Marine Environment Laboratory, IAEA, 4 Quai Antoine 1^{er}, MC-98000, Monaco (j.orr@iaea.org)

² Department of Chemistry, Göteborg University, Göteborg, S-412 96, Sweden

³LSCE, UMR CEA-CNRS, CEA Saclay, Gif-sur-Yvette, F-91191, France

CHANGES IN pH AND HYDROGRAPHIC PARAMETERS IN THE BALTIC SEA Perttilä, Matti

The pH, alkalinity, salinity and calcium values in the Baltic Sea are affected by human activities, hydrographic processes, primary production, global CO₂ increase and acid rain in the drainage area. Distinguishing the effects of these pressures is one of the aims of the Baltic Sea monitoring programme.

The partial pressure of CO₂ shows large seasonal and geographical variations in the surface layer of the Baltic Sea. During winter, in the ice-free part of the sea it is generally close to atmospheric partial pressure, part indicating a pH equilibrium. Average pH for the winter period in the central Baltic Sea surface water has decreased from about 8.2 in 1985 to present value 8.1. The average summer pH, however, is rising, probably indicating an increasing primary production. Similar behaviour is found in the Gulf of Finland.

Alkalinity and calcium data in the Baltic Sea are discussed in view of carbonate processes, salinity variations and fresh water inputs. The development of calcium – salinity ratios in both Gulf of Finland and Gulf of Bothnia indicate enhanced calcium dissolution in both drainage areas, leading to a [Ca]/TA ratio higher than expected by the simple mixing of North Sea high salinity water with fresh water.

The investigation on alkalinity, calcium and salinity emphasize the potential use of their relations as water mass tracers in the Baltic Sea, and as tracers of environmental changes especially in the low salinity regions.

Finnish Institute of Marine Research, PO Box 2, FIN-00561 Helsinki Finland

EFFECTS OF OCEAN ACIDIFICATION ON THE BACTERIAL DEGRADATION OF ORGANIC MATTER M39

Piontek, Judith, Nicole Händel, Mirko Lunau, Corinna Borchard, and Anja Engel

The amount of CO₂ being sequestered from the atmosphere to the ocean is mainly controlled by the balance between autotrophic production, calcification and microbial decomposition of organic matter. In recent studies, rising CO₂ was shown to reduce the calcification of marine phytoplankton and to affect the photosynthetic carbon acquisition of single phytoplankton species and natural phytoplankton communities. However, effects of ocean acidification on microbial degradation processes and potential consequences for the remineralization of carbon are not yet considered, despite the fact that the largest proportion of organic matter is degraded in the upper water column, the zone primarily affected by rising CO₂. We tested the effect of decreasing pH on the degradation of organic matter in a series of chemostat- and batch-experiments with cultures of Emiliania huxleyi and with coccolithophore-dominated natural phytoplankton communities as source of degradable organic matter. Results show that decreasing pH affects the degradation of organic matter as derived from changes in the activities of polysaccharide- and protein-degrading extracellular enzymes (e.g. glucosidases, leucine-aminopeptidase). Variations in the microbial degradation activity during our experiments were related to changes in the biogeochemistry of organic matter. Our findings demonstrate the necessity to include the response of marine bacteria to ocean acidification when estimating the future carbon cycle.

⁴Department of Biol. Sciences, Cal. State University San Marcos, San Marcos, CA, 92096-0001, U.S.A.

⁵Climate and Environmental Physics, University of Bern, Bern, CH-3012, Switzerland

⁶ Bedford Institute of Oceanography, Dartmouth, NS, B2Y 4A2, Canada

⁷ Max Planck Institut für Meteorologie, Hamburg, D-20146, Germany

⁸Université Catholique de Louvain, Institut d'Astronomie et de Geophysique Georges Lemaitre, Louvain-La-Neuve, B-1348, Belgium

Alfred Wegener Institute for Polar and Marine Research, Am Handelshafen 12, 27570 Bremerhaven, Germany (Judith.Piontek@awi.de)

MACROALGAL COMMUNITY RESPONSE TO INCREASING CO2

T39

Porzio, Lucia¹, Jason M. Hall-Spencer², and Maria Cristina Buia¹

By 2100 the surface waters of our oceans are predicted to have decreased in pH by 0.5 units due to anthropogenic emissions of CO₂ into the atmosphere (IPCC report, 2007). Many studies are now trying to find out how this will affect marine systems, mainly by using short-term mesocosm and laboratory studies.

Here we show how the community composition of shallow-sublittoral macroalgae varies along a natural pH gradient at Castello Aragonese on the island of Ischia (Italy) where volcanic vents emit 90-95% CO₂ (together with 3-6% N₂ and small amounts of O₂, Ar and CH₄) at ambient temperature. The venting CO₂ acidifies seawater along a gradient from 8.17 down to 6.57 along a 300 m transect running parallel to the rocky shore (Hall-Spencer *et al.*, 2008). We use this site to investigate which algae are tolerant of the long-term effects of increased CO₂ levels and how this affects macroalgal community structure.

Macroalgae were scraped from known areas of vertical rock faces to quantify the macroalgal community structure and their reproductive status. The most obvious effect was that both crustose and erect calcified algae (e.g. *Mesophyllum sp.*, *Neogoniolithon brassica-florida*, *Jania rubens* and *Padina pavonica*) first decreased their reproductive potential then disappeared from the community. These calcifiers were being replaced by other crustose and erect algae (e.g. *Hildenbrandia rubra*, *Dictyota dichotoma* and *Osmundea truncata*). Brown algae were the high-biomass canopy forming species throughout the CO₂ gradient but *Cystoseira amentacea* var. *stricta* was dominant at normal CO₂ levels whereas *Sargassum vulgare* became dominant in the acidified zone.

We noted a 25% reduction in macroalgal biodiversity in the acidified zone. Ecophysiological analysis are now needed to determine what is special about the highly tolerant species and whether calcified species can adapt their mineralogy to cope with small increases in CO₂ levels.

OCEAN ACIDIFICATION DUE TO MASSIVE CARBON DIOXIDE VENTING ALONG THE WAGNER FAULT, GULF OF CALIFORNIA, MEXICO. M40

<u>Prol-Ledesma, Rosa Ma.</u>¹, Carles Canet¹ and Paul R. Dando²

The Wagner Fault in the Northern Gulf of California, Mexico, is an area where CO₂ rises to the surface along deep faults in the 5 km-thick sedimentary column. There are an estimated 15, 000 gas vents in the area. The presence of bubbles, breaking the surface above the Wagner Fault vents, indicates CO₂ release to the atmosphere. Individual gas plumes can be 100 m or more across and bubbles reach the surface from 150 m depth resulting in acidification of the entire water column. Close to the outlets pH values can be as low as 6.3, with a full range of pH values up to 8.2 occurring in the water column and along the seabed. This makes the area a good study site for examining the effects of elevated CO₂ and low pH on the marine ecosystem. The gas venting has given rise to varied bottom topography with sorted, coarse-grained sediments and cemented deposits as well as the background silty-clays. This has resulted in a high diversity of bottom fauna, including groups with calcareous shells and skeletons. Since the water depths in the area do not exceed 210 m, these study sites are easily sampled using vessels with smaller ROVs.

¹Laboratorio di Ecologia del Benthos, Stazione Zoologica 'A. Dohrn', P.ta S. Pietro, 80077 Ischia (Naples), Italy. lucia.porzio@szn.it

²Marine Institute, Marine Biology and Ecology Research Centre, University of Plymouth, Plymouth PL4 8AA, U.K.

¹Departamento de Recursos Naturales, Instituto de Geofisica, Universidad Nacional Autónoma de México, Ciudad Universitaria, Delegación Coyoacán, 04510 Mexico D.F., Mexico

EFFECT OF ELEVATED PCO₂ ON THE BORON ISOTOPIC COMPOSITION AND B/CA RATIO INTO THE MEDITERRANEAN SCLERACTINIAN CORAL CLADOCORA CAESPITOSA T40

Reynaud, Stéphanie¹, Sophie Martin^{2, 3}, Claire Rollion-Bard⁴, Paolo Montagna⁵, Malcolm McCulloch⁶, Christine Ferrier-Pagès¹, and Jean-Pierre Gattuso²

It has been well established that ocean acidification negatively affects the calcification rate of calcifying organisms. However, how pH and the modified seawater chemistry act on the site of calcification is still debated. Because incorporation and fractionation of boron isotopes in carbonates are controlled by pH, boron is used as a proxy to understand coral calcification mechanisms.

The effect of the increase of CO_2 partial pressure (pCO_2) expected by 2100 was tested on the boron isotopic composition ($\delta^{11}B$) and the B/Ca ratio in the skeleton of the Mediterranean symbiotic coral, *Cladocora caespitosa*.

Colonies were collected in the Villefranche Bay (France) at 25 m depth. Corals were acclimated in aquaria under summer temperature (22°C) and irradiance (35 μ mol photon m⁻² s⁻¹) before being stained with alizarin. Colonies were split in two aquaria maintained at normal (400 ppm) and elevated pCO_2 (700 ppm). Temperature, light and photoperiod were changed according to natural yearly fluctuations at ca. 25 m depth in the Villefranche Bay. After one year of culture, the $\delta^{11}B$ and the B/Ca ratios were analyzed in the skeleton deposited above the alizarin line.

Preliminary results did not show any difference in $\delta^{11}B$ between the treatments: $27.3 \pm 1.4 \% vs. 27.2 \pm 2.3 \%$ respectively for 400 and 700 ppm. These values correspond to a pH of 8.75. The lack of difference between treatments has also been obtained with the B/Ca ratio $(0.926 \pm 0.063 \text{ and } 0.929 \pm 0.077 \text{ mmol/mol}$, respectively). We have concluded that on this range of seawater pH, the coral controls the pH at the calcification site.

LIGHT-DEPENDENT TRANSCRIPTIONAL REGULATION OF GENES OF BIOGEOCHEMICAL INTEREST IN THE DIPLOID AND HAPLOID LIFE STAGES OF EMILIANIA HUXLEYI M41

<u>Richier, Sophie</u>¹, Marie-Emmanuelle Kerros¹, Colomban de Vargas^{2, 3}, Liti Haramaty³, Paul G Falkowski^{3,4}, and Jean-Pierre Gattuso¹

The cosmopolitan coccolithophorid *Emiliania huxleyi* is capable of carbon uptake by both photosynthesis for organic matter production and by intracellular calcification for coccolith production. The algal cells, generating huge blooms, photosynthetically produce a large quantity of organic matter and calcite crystals by fixing dissolved inorganic carbons (DIC), before sinking to the bottom of the ocean. Consequently, the regulation of photosynthesis and calcification by coccolithophorids is a key factor in the global carbon budget and constitutes one of the main targets of high-CO₂ condition in the near future. In order to follow both photosynthesis and calcification process in *E. huxleyi* CCMP1516 haploid and diploid stages, we decided to focus on the ribulose-bisphosphate carboxylase-oxygenase (rbcL) gene as candidate for photosynthesis and the calcium binding protein (gpa) for calcification. Transcripts regulation was monitored using quantitative RT-PCR method on a

¹Centre Scientifique de Monaco, Av. Saint Martin, MC-98000 Monaco (sreynaud@centrescientifique.mc)

²CNRS-Université de Paris 6, Villefranche-sur-Mer, France

³Marine Environmental Laboratories, International Atomic Energy Agency, 4 Quai Antoine 1er MC-98012 Monaco

⁴CRPG-CNRS, Nancy-Université, 15 rue N.D. des Pauvres, Vandoeuvre-lès-Nancy, France

⁵ICRAM, Via di Casalotti 300, Rome, Italy

⁶RSES, ANU, Canberra 0200, Australia

diel cycle. A similar diel regulation of *rbcL* mRNAs, with a peak of expression 2 h after the beginning of the light period, was observed in both haploid and diploid cells. Similarly, *gpa* mRNAs were upregulated after 2 h of light but restricted to the haploid cells. However, *E. huxleyi* diploid cells, which precipitate calcium carbonate, exhibited a significantly larger abundance of *gpa* transcripts than the haploid, non-calcifiying cells, throughout the diel cycle. Comparative study has been conducted on *E. huxleyi* strain TQ26 and the analysis was extended to two more genes potentially involved in calcification such as the calmodulin and the carbonic anhydrase. The genes of interest are discussed as potential candidates for biogeochemical proxies.

RISING SATURATION HORIZONS MAY OVERTAKE COLD, DEEP CORALLINE COMMUNITIES IN ANTARCTICA T41

Riddle, Martin J.¹, Bronte Tilbrook^{2,3} Ian Snape, Kristina Paterson^{2,3}, and Katherine Berry³

A diverse, coral-based benthic community was recently discovered and recorded on video in a canyon system on the Antarctic continental slope off East Antarctica. The main structural components are heavily calcified branching hydrocorals and gorgonians creating habitat for the highest diversity of invertebrates and fish seen during a 20 day survey between 139°E and 145°E. The community was not seen at any of the ~60 shallower sites sampled on the continental shelf, possibly because it is slow growing and not competitive during early re-colonisation after ice-berg scour. The calcareous species were dominant at 600-850 m, well below the reach of modern ice-bergs, but were sparse at 900-1000 m and apparently absent at 1400 m. Water column CO₂ chemistry indicates that the aragonite saturation horizon was at about 900 m, with the calcite saturation state at this depth approximately The canyon system drains cold, saline Antarctic Bottom Water generated during sea-ice formation in the permanently open waters of the Mertz Polyna, west of the 90 km long Mertz Glacier. Katabatic winds from the continent accelerate ice formation in the polynya and are likely to increase air-sea exchange of CO₂. The continued uptake of CO₂ will drive lower the saturation state of carbonate minerals and we predict that these coralline benthic communities will be among the first globally to show calcification effects due to the crossing of saturation thresholds. The narrow depth range currently occupied by this community suggests its continued existence may be threatened if acidification further reduces its range.

COMPARISON OF HALIMEDA ARAGONITE FROM 40 YEARS OF ARCHIVED FIELD SAMPLES WITH SPECIMENS GROWN IN HIGH pCO $_2$ SEAWATER M42

Robbins, L.L.¹, P. Knorr^{1P}, P. Hallock², and K. Yates¹

Ocean chemistry is changing more dramatically now than in the last 20 million years because of rising atmospheric pCO₂ levels. In fact, pH values of the open ocean have decreased by 0.1 since 1980 and are predicted to decrease 0.3-0.5 in the next 80 years. Ocean acidification will likely affect fundamental geochemical and biological processes including calcification and carbonate sediment production. The west Florida shelf has a natural gradient from temperate to subtropical carbonate sedimentation which provides natural laboratories to examine the effects of ocean acidification on

¹Laboratoire d'Océanographie de Villefranche, Université Pierre et Marie Curie-Paris 6 / CNRS, 06234 Villefranche-sur-Mer Cedex, France (richier@obs-vlfr.fr).

² EPPO, UMR7144, CNRS – Station Biologique de Roscoff, F-29680 Roscoff, France

³Institute of Marine and Coastal Sciences, Rutgers The State University of New Jersey, New Brunswick, New Jersey, 08901, USA.

⁴Department of Geological Sciences, Rutgers, The State University of New Jersey, Piscataway, NJ 08854.

¹ Australian Antarctic Division, Channel Highway, Kingston, Tasmania 7050, Australia.

² Antarctic Climate and Ecosystems CRC, Hobart, Tasmania 7001

³ Centre for Climate and Weather Research, CSIRO, Hobart, Tasmania 7000, Australia.

aragonite production by calcareous green algae. Scanning electron microscopy (SEM) can reveal ultrastructural details of calcification that occurred at different saturation states. Comparison of *Halimeda* spp. from west Florida shelf archived samples collected more than 40 years ago, with specimens collected in ongoing studies, indicates the effect of increased pCO₂ and decreased saturation state over this time span. Laboratory experiments using living specimens of *Halimeda* in a range of pCO₂ environments are being used to constrain historical observations. *Halimeda* crystal data from apical sections indicate that increases in crystal density and decreased crystal width occurred over the last 40 years.

¹U.S. Geological Survey, St. Petersburg, FL 33701, USA (lrobbins@usgs.gov)
²University of South Florida, College of Marine Science, 140 7th Avenue South, St. Petersburg, FL 33701

PROJECTIONS OF VENTILATION IN A COUPLED CLIMATE MODEL: IMPLICATIONS FOR OCEANIC DISSOLVED OXYGEN T42

Russell, Joellen L. and Paul J. Goodman

Future circulation changes due to global warming are projected to have a dramatic impact on the concentration of dissolved oxygen in the global ocean. Warming of surface waters is likely to affect the concentration of dissolved oxygen in the ocean directly through solubility, as well as indirectly through stratification-driven changes to organic material export and remineralization rates. These changes are likely to lead to decreased oxygen in equatorial thermocline waters according to current climate model predictions. Results from the GFDL CM2.1 climate model indicate that the dissolved oxygen content of the Southern Ocean may increase temporarily during the transient into a high-CO2 world through increased wind-driven ventilation of cold, low-oxygen waters. Contrary to previous work, increased wind-forced upwelling associated with the poleward intensification of the Southern Hemisphere Westerlies leads to increases in Southern Ocean dissolved oxygen concentration over the next 100 years, not the decreases seen in models with poorly resolved Southern Hemisphere Westerly Winds.

Department of Geosciences, University of Arizona, 1040 E 4th St., Tucson, AZ 85719, USA (jrussell@email.arizona.edu)

UPWELLING OF 'ACIDIFIED' WATER ONTO THE NORTH AMERICAN CONTINENTAL SHELF M43

Sabine, Christopher L.¹, Richard A. Feely¹, J. Martin Hernandez-Ayon², Debby Ianson³, and Burke Hales⁴

The absorption of atmospheric carbon dioxide into the ocean lowers the pH of the waters. This so-called ocean acidification could have important consequences for marine ecosystems. In order to better understand the extent of this ocean acidification in coastal waters, we surveyed 13 hydrographic sections perpendicular to the coast between Queen Charlotte Sound, Canada and the Mexican Baja Peninsula between May and June of 2007. Upwelling was observed from line 2 off central Vancouver Island to line 13 off Baja California, Mexico. Lines 2 and 3 to the north and lines 8 to 11 to the south experienced the weakest upwelling. Recent upwelling was observed on lines 5 and 6 near the Oregon-California border. Coincident with the upwelled waters we observed low pH seawater that was undersaturated with respect to aragonite covering large portions of the continental shelf. The undersaturated water reached depths of approximately 40-120 m along most transect lines and all the way to the surface on line 5. The aragonite saturation horizon closely followed the 26.2 potential density surface where the waters had a pH of about 7.75. This density surface shoaled from a depth of ~175 m in the offshore waters and breached the surface over the shelf near the 100 m bottom contour, approximately 40 km from the coast. While seasonal upwelling of the undersaturated waters onto the

shelf is a natural phenomenon in this region, the ocean uptake of anthropogenic CO₂ has increased the areal extent of the affected area.

¹Pacific Marine Environmental Laboratory/NOAA, 7600 Sand Point Way NE, Seattle, WA 98115-6349. USA (chris.sabine@noaa.gov)

²Instituto de Investigaciones Oceanologicas. Universidad Autonoma de Baja California. Km. 103 Carr. Tijuana-Ensenada. Ensenada. Baja California. Mexico.

³Fisheries and Oceans Canada, Institute of Ocean Science, P.O. Box 6000, Sidney, BC V8L 4B2, Canada

⁴College of Oceanic & Atmospheric Sciences, Oregon State University, 104 Ocean Admin. Bldg., Corvallis, OR 97331-5503, USA

ACIDIFICATION SENSITIVITY OF THE BLUE MUSSEL, MYTILUS EDULIS, IN THE **BALTIC SEA: A HOLISTIC APPROACH**

Saphörster, Julia, Jörn Thomsen, Agnes Heinemann, and Frank Melzner

Blue mussels of the genus Mytilus from the North Sea and the Mediterranean displayed decreased rates of calcification in combination with metabolic depression and uncompensated extracellular pH when exposed to low ocean pH. Baltic Sea M. edulis might even react more sensitively to acidified waters than populations from full-strength saline oceans, as the dilute Baltic is characterized by a low buffering capacity (TA <2.0 meq/l) and low calcium carbonate saturation (Ω_{arag} <1.5). Employing a holistic approach, we related acid base parameters, ion exchange capacity and extracellular ion composition to whole-animal performance indicators and shell isotopic composition.

Adult mussels were kept in flow through aquaria for 14 days at five different CO₂ concentrations ranging from 380 ppm to 1,400 ppm. During the experimental period, we determined heart rates and valve/siphon opening status at the lowest and highest CO₂ levels as measures for overall organismic performance. Following the incubation period, hemolymph samples were drawn from the posterior shell retractor muscle of all animals for extracellular pH measurements. Tissue samples were taken for determinations of intracellular pH and Na⁺/K⁺ ATPase activity. In addition, ion composition of hemolymph and extra-pallial fluid will be investigated by means of ion chromatography. All biochemical / physiological parameters will be correlated with whole animal performance indicators to gain a comprehensive understanding of the potential vulnerability of the species towards future ocean acidification. Since Mytilus edulis is one of the most important ecosystem engineers in northern seas, its acidification sensitivity might have significant ecological consequences.

IFM-GEOMAR, Leibniz Institute of Marine Sciences (at Kiel University), Hohenbergstr. 2, 24105 Kiel, Germany (jsaphoerster@ifm-geomar.de)

ANTHROPOGENIC GAS DISTRIBUTION IN THE SOUTHERN OCEAN: FORMATION **AND SPREADING OF SAMW AND AAIW**Sasai, Yoshikazu ^{1,2}, Akio Ishida ^{1,2,3}, Yasuhiro Yamanaka ^{1,2,4}, and Hideharu Sasaki ⁵

Chemical tracers are widely used to investigate the ocean circulation, water mass formation rate, and mixing and ventilation processes. Anthropogenic gas such as chlorofluorocarbon (CFC) is inert transit tracer in the ocean, similar in some ways to anthropogenic CO₂, and provides information on the ocean circulation and its variability on timescale of months to decade. We have carried out CFC-11 simulation to investigate the formation and spreading of Subantarctic mode water (SAMW) and Antarctic intermediate water (AAIW) using an eddy-resolving ocean general circulation model. The model has a horizontal grid scale of 0.1 degree and 54 vertical levels. A model is forced by a climatological data of NCEP/NCAR. The model exhibits considerable skill in reproducing the section along Greenwich Meridian (AJAX), 30E (WOCE I06SB), 115E (WOCE I09S), and 170W (WOCE P15S). The high CFC-11 water penetrates to 1000 m depth between 50S and 40S by the SAMW and AAIW ventilation for each section. In the distribution of CFC-11 inventory in 1994, the model is a good performance to observed distribution (GLODAP), especially between 60S and 30S. Distribution of CFC-11 inventory in 26.5-27.0 (SAMW) and 27.0-27.4 (AAIW) in the model is also close to observed. The eddy-resolving model explicitly simulates meso-scale eddies and improves the structure of upper ocean stratification in the Southern Ocean.

¹ Frontier Research Center for Global Change, Japan Agency for Marine-Earth Science and Technology, Yokohama, 236-0001, Japan (ysasai@jamstec.go.jp)

² Core Research for Evolutional Science and Technology (CREST), Japan Science and Technology Agency (JST), Kawaguchi, 332-0012, Japan

³ Institute of Observational Research for Global Change, Japan Agency for Marine-Earth Science and Technology, Yokosuka, 237-0061, Japan

⁴ Graduate School of Environmental Earth Science, Hokkaido University, Sapporo, 060-0810, Japan

⁵ Earth Simulator Center, Japan Agency for Marine-Earth Science and Technology, Yokohama, 236-001, Japan

EFFECTS OF HYPERCAPNIA AND HYPOXIA ON THE EMBRYONIC DEVELOPMENT OF RED SEA BREAM, *PAGRUS MAJOR* T44

Sawada, Yoshifumi¹, Kazuhiro Higuchi², Tomoki Honryo³, Manabu Seoka⁴, Tokihiko Okada⁵, Yutaka Haga⁶, and Kazuhiro Ura⁷

Because of the weak stress resistance of early stage aquatic animals, hypercapnic and hypoxic conditions are supposed to cause a mass mortality and/or teratogenic effects leading to a large adverse effect on their population dynamics. However, most of the effects of these conditions on their early development are less well understood. This paper demonstrates the effects of hypercapnia and hypoxia on the embryonic development of a teleost red sea bream (RSB), *Pagrus major*, which is an important constituent of neritic ecosystem and an important fisheries resource in the Far East.

RSB eggs were exposed to seawater of six dissolved oxygen concentrations (0, 10, 25, 50, 75, and 100%) and four dissolved carbon dioxide concentrations (0, 60, and 120 mg/L) for seven different periods (0-180 min.) during somitogenesis. Larvae hatched from these eggs were observed the condition of somites by microscopy. Isolated exposure to hypercapnia and hypoxia induced somitic disturbances at high incidence rates in newly hatched larvae. Combined condition of hypercapnia and hypoxia induced somitic disturbances in RSB larvae at the higher incidence rates than the isolated ones.

Our previous study demonstrated that the somitic disturbances in newly hatched larvae develop into centrum defects in the vertebral bone. The results of this study suggest that hypoxic and hypercapnic conditions in aquatic environment caused by the human-induced excess organic matter loading, red tides, and the planned carbon dioxide disposal into the sea are the possible causes of the reduction of wild fish population by the induction of developmental defects.

¹Professor, Fisheries Laboratory, Kinki University, Ohshima 1790-4, Kushimoto, Wakayama 649-3633, Japan (YoshifumiSawada@za.ztv.ne.jp)

²Graduate student, ³ Engineer, ⁴ Assistant professor ⁵ Chief engineer, Fisheries Laboratory, Kinki University

⁶Assistant professor, Faculty of Marine Science, Tokyo University of Marine Science and Technology, Konan 457, Tokyo 108-8477, Japan

⁷Assistant professor, Graduate School of Fisheries Science, Hokkaido University, Hakodate, Hokkaido 041-8611, Japan

PAST ANALOGUES FOR OCEAN ACIDIFICATON: INSIGHTS FROM THE PALEOCENE/EOCENE THERMAL MAXIMUM M45

Schmidt, D.N.¹, A. Ridgwell², S.A. Kasemann³, and E.Thomas⁴

Continued absorption of fossil fuel CO_2 by the ocean may decrease surface ocean pH by up to 0.5-0.6 pH units over the next 250 years, pushing ocean geochemistry outside the range of values experienced during the last tens of millions of years. The Palaeocene-Eocene Thermal Maximum (PETM) has been suggested as an analogue for future global warming and ocean acidification, but pH and carbonate system changes at this time have not been well constrained.

We employed high-spatial resolution secondary ionization mass spectrometry (SIMS) of tests of the benthic foraminifer *Oridorsalis umbonatus* at Maud Rise (Site 690B, Weddell Sea, Southern Ocean, palaeodepth 1900 m) and *Lenticulina* sp. at Bass River (Leg 174X, New Jersey; shelf) to characterize B/Ca (carbonate saturation proxy) and Li/Ca (DIC proxy). Mg/Ca data indicate a total temperature increase from 12.7°C to 18.5°C at Maud Rise, in agreement with previous work, and a 4°C temperature increase at Bass River to 36°C during the PETM. The latter needs further evaluation because Mg/Ca for *Lenticulina* has not been calibrated. B/Ca show a transition from slightly oversaturated to under-saturated conditions going into the PETM δ^{13} C excursion at both sites and do not recover to pre-event values, even after recovery of δ^{13} C and temperature values. Li/Ca ratios show a minor reduction at both Sites, indicating little change in DIC, in contrast to what is expected in view of the magnitude of carbon release; we need more information on the behaviour of this proxy. Ba/Ca indicate an increase in productivity at Bass River but a decrease at Maud Rise, in agreement with micropalaeontological evidence.

² School of Geographical Sciences, University of Bristol, Bristol, BS8 1SS, U.K.

SENSITIVITY OF LYSOCLINE DEPTH AND AIR-SEA CO₂-EXCHANGE TO PARTICLE FLUXES IN THE WATER COLUMN T45

Schneider, Birgit^{1,2}, Laurent Bopp¹, and Marion Gehlen¹

An ocean biogeochemical model was applied to estimate the sensitivity of lysocline depth and air-sea CO₂-exchange to the vertical flux of particulate organic and inorganic carbon (POC, PIC). The results of four different perturbation experiments show that instaneously induced changes in the sinking of POC and PIC cause large scale reorganizations in the marine ecosystem that also result in shifts of the lysocline depth after 100 years of model integration. Although similar effects with a general deepening of the lysocline in the Atlantic and the Southern Ocean next to a shoaling in the North Pacific are observed in all simulations, the re-equilibration of air-sea CO₂-fluxes yields oceanic DIC inventories that are either higher or lower than in the reference simulation. Based on fluxes of POC and PIC, productivity and the distributions of nitrogen (NO₃), dissolved inorganic carbon (DIC) and alkalinity a number of indices defined to estimate the efficiency of carbon transport away from the atmosphere. The results show that the more efficient the vertical transport of organic carbon towards depth, the lower the surface ocean pCO₂, the higher the air-sea CO₂ flux and the stronger the increase in the oceanic inventory of dissolved in organic carbon (DIC). Along with POC flux it is important to consider variations in the flux of PIC, as the net effect of particle flux reorganizations on surface ocean pCO₂ is a combination of changes in DIC and alkalinity.

¹ Department of Earth Science, University of Bristol, Bristol, BS8 1RJ, U.K.; d.schmidt@bristol.ac.uk

³ Grant Institute of Earth Science, The University of Edinburgh, The King's Buildings, Edinburgh, EH9 3JW, U.K.

⁴Department of Geology and Geophysics, Yale University, New Haven, CT 06520-8109, USA

¹ Laboratoire des Sciences du Climat et de l'Environnement (LSCE), F-91191 Gif-sur-Yvette, France.

² now at: Institute of Geosciences, University of Kiel, D-24098 Kiel, Germany.

COUPLED PHYSICAL BIOGEOCHEMICAL RESPONSE TO INCREASING CO₂ FROM AN EARTH SYSTEM MODEL M46

Segschneider, Joachim and E. Maier-Reimer

The recently intensified discussion about the impact of increasing uptake of CO_2 by the oceans and the subsequent acidification on the marine biota has mainly been focusing on the biogeochemical system. Here, by using results from a fully coupled Earth System Model driven with A2 CO_2 emissions, it will be shown that the physical response of the ocean to climate change induced by increasing atmospheric CO_2 may be of equal importance as the biogeochemistry:

The slowdown of the meridional overturning circulation in response to climate warming causes reduced upwelling of nutrients. This effect is more pronounced for silicate than for nitrate, since biogenic opal remineralizes at greater depth than particulate organic carbon. Hence, silicate concentrations in the surface waters decrease more rapidly than for phosphate and nitrate. Diatoms, which currently outcompete coccolithophores in the request for nitrate and phosphate provided that sufficient silicate is available, are thus having a relative disadvantage. In our model study, this results in an increase of global calcium carbonate export of 20% by the year 2100. Since this is quite contrary to what one would expect from a purely biogeochemical view (namely the decrease of calcite formation in more acid waters) it should be noted that our model computes enhanced dissolution of calcium carbonate (with an predicted drop in pH of about 0.3 units in surface water, which are still supersaturated), but no impact of acidification on the calcite formation is included.

Max-Planck-Institute for Meteorology, Bundesstr. 53, D-20146 Hamburg, Germany (joachim.segschneider@zmaw.de)

DISSOLVED INORGANIC CARBON IN THE CANADIAN ARCHIPELAGO OF THE ARCTIC OCEAN: THE EXPORT OF PACIFIC CARBON TO THE NORTH ATLANTIC VIA BAFFIN BAY T46

Shadwick, E.H., H. Thomas, A. E.F. Prowe, S. Moore, and D. Leong

The Arctic Ocean appears to be particularly sensitive to climatic changes, and is thought to be an area where such changes might be detected. The Arctic hydrological cycle is influenced by: runoff and precipitation, sea ice formation/melting, and the inflow of saline waters from the Bering and Fram Strait and the Barents Sea Shelf. Pacific water is recognizable as low salinity water, with high concentrations of dissolved inorganic carbon (DIC), flowing from the Arctic Ocean to the North Atlantic via the Canadian Archipelago. We present DIC data from an east-west section through the Archipelago, as part of the Canadian IPY initiatives. The fractions of Pacific and Arctic Ocean waters leaving the Archipelago and entering Baffin Bay and subsequently the North Atlantic are computed. The transport of carbon from the Pacific, via the Arctic, to the North Atlantic is estimated.

Fluctuations in water mass transport out off the Arctic have been linked to the North Atlantic Oscillation (NAO). During NAO induced Great Salinity Anomalies (GSA), large amounts of fresher water, with relatively higher concentrations of DIC, are released to the North Atlantic. The contribution of Pacific-origin carbon exported to the North Atlantic during such an anomaly is estimated. The reorganization of the Arctic circulation during the recent decades has led to an altered transport of Pacific waters to the North Atlantic Ocean. This in turn is deemed to modulate CO₂ uptake and acidification in the temperate and subpolar North Atlantic.

Dept. of Oceanography, Dalhousie University, Halifax, NS, Canada (Elizabeth.shadwick@dal.ca)

EFFECT OF THE HIGH CO₂ ON GROWTH OF PHYTOPLANKTON SPECIES: LABORATORY STUDY M47

Shin, Kyoungsoon, Bong-Gil Hyun, and Kitack Lee

Cultured phytoplankton, *Cylindrotheca closterium*, *Melosira nummuloides*, *Nitzschia* sp., *Phaeodictylum triconutum*, *Prorocentrum minimum* were subjected to test the growth rates under the different conditions of CO_2 concentration. Two different CO_2 concentrations were defined as 'normal condition' (p CO_2 : 200ppm) and 'high condition' (p CO_2 : 980ppm). The experiment was performed under the nutrient enriched-condition. The temperature was maintained 25 ± 1 under the equally lit light cycle (12 hr. light: 12 hr. dark). *Nitzschia* sp. and *M. nummuloides* showed higher chlorophylla concentration under the high condition than under the normal condition. However, higher chlorophylla concentration was measured for *P. minimum* under the normal condition than under the high condition. These results may reflect that the highly concentrated CO_2 environment increased the growth rate of *Nitzschia* sp. and *M. nummuloides*, while the growth rate of *P. minimum* was retarded under the high condition. There was no a clear difference of growth rate between high and normal conditions in the case of *C. closterium* and *P. triconutum*. The researches are in progress and will argue on the matter of the rapid variations of phytoplankton community structure and primary productivity which is induced by the increase of global CO_2 level.

NATURAL ANALOGUE FOR OCEAN ACIDIFICATION Shitashima, Kiminori T47

Large amounts of CO2 are being supplied to the ocean from the seafloor as a natural phenomenon at seafloor hydrothermal systems. Hydrothermal fluids are highly enriched with CO2 and show low pH (around pH 2 to 3) relative to seawater. This CO2 is taken up from the basalt by the fluid during high temperature seawater-magma interaction. In the Okinawa Trough and Mariana Trough, liquid CO2 is emitted from hydrothermal vents at about 1500m depth, and CO2 gas bubbles are erupted from seafloor at 100-200m depth in the Kagoshima Bay.

Dissolution of CO2 during ascent of CO2 droplet and diffusion of low pH seawater were observed at the Hatoma Knoll in the Okinawa Trough. The CO2 droplets emitted from the seafloor dissolve slowly into the ambient seawater while ascending, but changes in pH and pCO2 near the rising CO2 droplets are small. The in-situ pH mapping revealed that the discharged liquid CO2 does not cause widespread pH depression in the ambient environment. The result of CO2 gas bubbles mapping survey at the Wakamiko Caldera in the Kagoshima Bay showed only localized pH depression below 120m depth. At the NW Eifuku submarine volcano in the Mariana Trough, the low pH plume derived from hydrothermal liquid CO2 was detected in 100m high and 200m wide area above the summit of the volcano.

Natural analogue of the hydrothermal liquid CO2 would provide an opportunity for understanding the mechanism, influence and recovery of ocean acidification.

Central Research Institute of Electric Power Industry, 1646 Abiko, Abiko-City, Chiba, 270-1194, JAPAN (shita@criepi.denken.or.jp)

¹Korea Ocean Research and Development Institute/South Sea Institute, Jangmok, 656-830, Korea (ksshin@kordi.re.kr)

²Pohang University of Science and Technology, School of Environmental Science and Engineering, Pohang, 790-784, Korea

EFFECTS OF OCEAN ACIDIFICATION ON DEVELOPMENT AND GROWTH OF S. DROEBACHIENSIS LARVAE AND ADULTS M48

Stumpp, Meike, Katja Trübenbach, and Frank Melzner

Previous work has shown that larval stages of sea urchins react highly sensitive towards ocean acidification, displaying dramatically reduced growth rates. However, nothing is known so far about the underlying physiological mechanisms leading to growth depression. Within the current study, we use the sea urchin *Strongylocentrotus droebachiensis*, a temperate to polar sea urchin that occurs commonly in all northern regions of the world oceans, as a model organism to study the effects of ocean acidification and global warming on key physiological processes.

Currently, basic experiments estimate the effects these environmental changes have on the development, morphology and survival of *S. droebachiensis* larvae and on the growth rate of adults. First results showed a significant reduction in larval growth rates at 1,400 ppm CO₂. Using quantitative PCR and the sequenced genome of the congener *S. purpuratus*, we will subsequently target gene products responsible for ion homeostasis, pH regulation and calcification, as these might be processes most relevant for the observed changes in larval morphology. Furthermore we will define tissue specific gene expressions responsible for ion homeostasis in adults. Additional non-hypothesis driven approaches, employing 454 pyrosequencing of control vs. stressed phenotype transcriptomes, will help us develop new hypotheses regarding what physiological processes might define sensitivity towards elevated CO₂ and temperature.

IFM-GEOMAR, Leibniz Institute of Marine Sciences (at Kiel University), Hohenberg Str. 2, 24105 Kiel, Germany (mstumpp@ifm-geomar.de)

REGULATION OF PHOTOSYNTHESIS OF A COCCOLITHOPHORID LIVING IN A HIGH CO2 WORLD T48

Suggett, David J., Richard Webster, Tracy Lawson, Stephane Lefebvre, Christine Raines, and Richard J. Geider

Growth of calcifying unicellular coccolithophorids contributes significantly to export of both organic carbon and calcium carbonate (CaCO₃) from surface waters. Previous research has demonstrated that calcification by these organisms will be negatively affected as the pH of the oceans decreases with rising atmospheric CO₂; however, the simultaneous response of photosynthesis and therefore growth is less clear. As with these previous studies we examined the globally important coccolithophorid Emiliania Huxleyi. A calcifying strain (PML-B11) was grown in pH stats at each of two CO₂ (180 and 750 ppm) and light (30 and 300 µmol photons m-2 s-1) growth conditions. In addition to calcification, we examined the response of the light harvesting antennae, photosynthetic reaction centers and photosynthesis rates (electron transfer rates, gross and net O₂ evolution, respiration). Calcification rates changed with CO2; however, most photosynthetic parameters were largely altered by light availability and not CO2. Only cell size significantly increased at higher CO2 levels. Similarly, light and not CO2 altered the expression of a subset of FCP genes at the RNA and protein levels. Our results further suggest that acidification will have limited effect on the photosynthetic capacity of coccoltihophorids but that combing calcification and cell size may have important ecological and biogeochemical implications. Our results are somewhat inconsistent with other similar studies highlighting that such fundamental responses must be better placed within the context of the genetic diversity for *E. huxleyi* that exists in nature.

Department of Biological Sciences, University of Essex, Colchester, CO4 3SQ, U.K.

A NEW APPROACH TO STUDY THE EFFECTS OF CO₂ IN DEEP-SEA ORGANSIMS M49 Thatje, Sven¹, Olaf Heilmayer¹, Bruce Shillito², Christian Osseforth¹, and Nelia Mestre¹

The deep ocean is one of the last of the planet's major frontiers. The ocean floor beyond the edge of the continental shelf covers some 50 percent or more of the entire Earth's surface. Deep-sea biology has traditionally been rather descriptive and dependent on the availability of organisms obtained by means of trawls or grabs. Research over the past decades has revealed high levels of biodiversity and endemism associated with many deep-sea ecosystems. Global warming and associated euthropication will have tremendous impact on deep-sea faunas that are assumed to be highly sensitive to changes in pH and temperature. Increased acidosis of the deep sea may lead to metabolic suppression, reduced protein synthesis, respiratory stress, reduced metabolic scope, and ultimately death of the organisms exposed to it. Only in recent years the increased use of underwater video systems and ROVs has allowed in situ studies of deep-sea organisms in their environment. Here, we advocate that the study of deep-sea organisms under controlled lab conditions (monitored experimental parameters) is still essential to elucidating ecological and physiological life history adaptations of deep-sea organisms to their environment, complementing field observations. We present an infrastructure for work on living deep-sea organisms under controlled laboratory conditions: The IPOCAMP_(TM) system, allows the study of organisms under deep-sea conditions including controlled conditions of temperature, pressure and carbon dioxide, in a closed seawater circulation system. We present first results of this newly established system in the CO₂ context.

CO₂-DRIVEN ACIDIFICATION RADICALLY AFFECTS LARVAL SURVIVAL AND DEVELOPMENT IN MARINE ORGANISMS – II. ECHINODERMS T49

Thorndyke, Michael¹, Jon Havenhand², and Sam Dupont¹

Perhaps one of the key marine groups most likely to be impacted by predicted OA is the echinoderms. Echinoderms are a vital component of the marine environment with representatives in virtually every ecosystem; where they are often keystone ecosystem engineers. In addition, many are indirect developers (e.g. urchins, brittlestars) where both larva and adult have critical (and quite different) episodes of skeletogenic calcification. In contrast others (Asteroids) exhibit adult but not larval skeletogenesis. In this way echinoderms offer a valuable and tractable experimental system for exploring the impacts of OA on marine biota. We have used our CO₂ – based sea water acidification system to investigate the affects of increasing acidity on early developmental success in the brittlestars Amphiura filiformis and Ophiothrix fragilis, the sea star Asterias rubens and the sea urchin Strongylocentrotus droebachiensis. Our data show that the impact of acidification on larval development is species-specific and highlight the danger of extrapolation from just a few key species, even within closely related taxa. For example, in O. fragilis, low pH induces dramatic mortality after one week due to larval and skeletal malformations while more subtle effects were observed in A. filiformis where some individuals survive to the juvenile stage. Urchins and asteroids on the other hand seem relatively unaffected. These results are interpreted in the context of adaptive potential to these rapid environmental changes.

¹ National Oceanography Centre, Southampton, School of Ocean and Earth Science, European Way, SO14 3ZH Southampton, U.K. (svth@noc.soton.ac.uk)

² UMR CNRS 7138, Systématique, Adaptation et Evolution, Université Pierre et Marie Curie, 7 Quai Saint Bernard, Batiment A, 4ème, 75252 Paris Cedex 05, France

¹Sven Lovén Centre for Marine Sciences - Kristineberg, 45034 Fiskebäckskil, Sweden

²Sven Lovén Centre for Marine Sciences - Tjärnö, 45296 Tjärnö, Sweden

THE MEDITERRANEAN SEA: A VERY HIGH ANTHROPOGENIC CO₂ AND ACIDIFIED ENVIRONMENT. M50

Touratier, F. and C. Goyet.

The Mediterranean Sea ecosystems suffer from a very intense anthropogenic pressure that strongly affects most biogeochemical cycles. During the last century, the impact on the carbon cycle and especially on the carbonate system has been poorly understood given the scarce data available for total alkalinity (A_T) , total dissolved inorganic (C_T) , and/or pH. Since almost a decade however, several national and international programs were designed to increase the amount of high quality data in the region.

Using this recent database we first describe the distributions for C_T and A_T for the whole Mediterranean Sea. The concentration of anthropogenic CO_2 (C_{ant}) accumulated since the preindustrial era and consequently the level of acidification are also estimated. It is shown that all waters (even the deepest ones in both the western and eastern basins) are contaminated by C_{ant} . This trend is confirmed by the distribution of other anthropogenic tracers like CFCs and tritium. We show that the C_{ant} averaged inventories in the Mediterranean Sea are approximately twice as those estimated in the North Atlantic Ocean, resulting in a much larger acidification of this semi-enclosed sea. Consequently, at depths, the Mediterranean Sea acts as a significant source of anthropogenic carbon for the Atlantic Ocean.

The analysis of the temporal trends clearly reveals the impact of the EMT (Eastern Mediterranean Transient) which resulted in an intense sequestration of anthropogenic carbon into the deep waters of the eastern basin. The recent observation that C_{ant} in the intermediate and deep waters of the DYFAMED site (western Mediterranean) is decreasing since the 1990's is also discussed in the light of the EMT event.

IMAGES, University of Perpignan Via Domitia, 56 av. Paul alduy, 66000 Perpignan, France.

THE EFFECTS OF ELEVATED CO₂ AND REDUCED pH ON THE INTERTIDAL SEA ANEMONE ANTHOPLEURA ELEGANTISSIMA AND ITS ALGAL SYMBIONTS Towanda, Trisha and Erik V. Thuesen

Anthopleura elegantissima (Brandt) is a non-calcifying anthozoan with symbiotic algae similar to that of hermatypic corals. It is locally abundant in the intertidal zone along the Pacific coast of North America. To better understand the impacts of ocean acidification on photosynthetic symbioses, the effects of elevated carbon dioxide (hypercapnia) and reduced pH were investigated on A. elegantissima. The anemone forms aggregations of clones through fission, and experiments were designed to examine effects of hypercapnic acidification with paired experiments using couplets of clones. Couplet individuals were maintained in the laboratory at pH values of 8.1 (ambient) or 7.3 (CO₂-induced) for a period of 10 weeks. Individuals were compared for differences in respiratory rate, photosynthetic rate, growth rate and the contribution of zooxanthellae to the animal's respiration (CZAR). Chlorophyll content and density of algal cells in anemones as well as algal cell size and mitotic index were also measured for comparisons.

Evergreen State College, Laboratory I, Olympia, WA 98505, USA (trisha towanda@yahoo.com)

TESTING THE SENSITIVITY OF THE CARBON CYCLE IN THE EASTERN MEDITERRANEAN SEA IN RESPONSE TO ANTHROPOGENIC INFLUENCE. M51 Triantafyllou, G., V. Avgoustidi, E. Krasakopoulou, K. Tsiaras, and A. Pollani

A seawater carbon cycle model adopted from Chuck et al (2005) has been implemented in the area of the Eastern Mediterranean Sea in an attempt to test the behavior of an ultra oligotrophic system, in view of the ongoing ecosystem changes. More specifically, we are aiming to asses the behavior of the seawater carbon cycle in the area under investigation, through the response of a number of biotic and

abiotic processes to climate change and anthropogenic influence. The studied area is considered to be highly sensitive to climate change and forms a system characterized by extreme phosphorous limitation all year round.

The model describes phytoplankton standing stocks and growth in the euphotic zone (120 m), seawater concentrations as well as riverine inputs of nitrate, phosphate and silicate, TCO₂, alkalinity, the air sea gas exchange of CO2, calcite production and remineralisation. Phytoplankton and phosphorous dynamics are provided by the ERSEM (European regional seas ecosystem model) in the carbon cycle model. Sensitivity experiments are being carried out through a series of perturbation tests; increased usage of fertilizers, increasing temperature, increased air sea gas exchange of CO₂ according to a number of different emissions scenarios, changes in the abundance of calcite/coccolithophores, changes in the action of the biological pump.

Such sensitivity analysis in the area of the Eastern Mediterranean Sea will allow a first investigation on the possible biogeochemical feedbacks that might arise in response to the current anthropogenic forcing.

Hellenic Centre for Marine Research, Anavissos, Greece. (gt@ath.hcmr.gr)

DISTRIBUTION OF PELAGIC BIOGENIC CARBONATES ACROSS THE SOUTHERN OCEAN SOUTH OF AUSTRALIA T51

Trull, Tom¹, Tim Smit², and Diana Davies¹

Developing an understanding of the impact of ocean acidification on pelagic microbial biogenic carbonate forming organisms requires knowledge of their abundance. As a first step in this assessment for Southern Ocean waters south of Australia, surface water PIC was measured from March 22, 2008 through April 19, 2008 at 60 sites along the WOCE/CLIVAR SR3 transect between Tasmania and Antarctica. PIC was determined on bulk samples collected from the underway clean seawater supply with a continuous flow centrifuge size-fractionated samples obtained by sequential filtration through a 1000 um pre-screen followed by 200 and 55 um nylon screens, and 1 um glassfibre filters. Bulk PIC values ranged between <1 ug/L and 20 ug/L. PIC values were low (1-2 ug/L) close to the Antarctic continent and increased (2-6 ug/L) in the AZ (Antarctic Zone) and PFZ (Polar Frontal Zone). Highest values were observed across the SAZ (Subantarctic Zone) with a peak of 16-20 ug/L between 51°S and 47°S. Results from the size-fractionated PIC measurements suggest that coccolithophores (the 1-55 um size fraction) made up the largest fraction (65-95%), with foraminifera (55-200 um fraction) contributing 15-30%, and the remaining small fraction of PIC came from other larger calcifying organisms like pteropods (200-1000 um fraction). The PIC distributions will be compared to carbonate saturation indices, and to particulate organic carbon and particulate biogenic silica distributions to provide a perspective on the relative importance of these carbonate forming taxa in comparison to other phytoplankton functional groups.

1. CSIRO Marine and Atmospheric Research, University of Tasmania, Antarctic Climate and Ecosystems Cooperative Research Centre, Hobart, 7001, Australia, Tom.Trull@utas.edu.au 2. Utrecht University, Netherlands

RESPONSES OF MARINE COMMUNITIES TO PH CHANGE. A FIELD BASED EXPERIMENTAL APPROACH. M52

Vance, Thomas

Without field based experimentation, accurately gauging the effect of increasing atmospheric pCO₂ on multi-species community processes such as recruitment, succession, trophic interactions, biogeochemical cycling and ecosystem functioning is challenging.

Obtaining such data is difficult in the lab, and due to the inherent pH buffering potential of seawater, has largely been rejected as not practical for a field based experimental approach.

This current study presents a novel field based method where settlement panels are exposed to experimentally manipulated CO₂ induced reduced pH environments. The pH conditions are manipulated to simulate future global ocean carbonate states. Microbial, algal and invertebrate communities were measured to provide benthic ecosystem responses. This approach allows a balanced and replicated field based experimental approach to be applied to answer fundamental questions about biological responses of communities to ocean acidification.

University of Newcastle upon Tyne, Biology, 5th Floor Ridley Building, Claremont Road, Newcastle upon Tyne, NE1 7RU, U.K. (thomas.vance@newcastle.ac.uk)

DECREASING OF AGGREGATE SIZE OF MARINE SUSPENDED PARTICLES UNDER HIGH CO₂ CONDITIONS T52

Watanabe, Yuji^{1,2}, Nobuhiro Maeda^{1,2}, and Koh Harada³

Suspended particles flocculate and form larger sinking particles in the ocean. Such large particles have an important role on the biological pump transporting organic carbon from the surface ocean to the deep layer, and the aggregates also provide the suitable habitat for bacteria and nano-flagellates. As the process of aggregation is affected by the surrounding pH conditions, we investigated the aggregation ability of marine particles by exposing them to lowered pH controlled with elevated CO₂. Suspended particles, which were collected from near the sea bottom (100 m depth) in Toyama bay and from about 1500 m depth in the western north Pacific, were stirred in seawater with various pH (6.0 - 8.0) conditions. The larger aggregates were formed in normal pH condition, but lowered pH condition inhibited the aggregation of suspended particles. Such decreasing the aggregate size causes decreasing in the sinking rate of that, consequently the available abundance of food source for planktonic organisms will be affected. And also the microhabitat for the attached microorganisms will decrease.

¹The General Environmental Technos Co., Ltd., Azuchimati, Chuo-ku, Osaka, 541-0052, JAPAN (watanabe yuji@kanso.co.jp)

²Research Institute of Innovative Technology for the Earth, Kizugawadai, Kizugawa, 619-0292, JAPAN

³Research Institute for Environmental Management Technology, National Institute of Advanced Industrial Science and Technology, Onogawa, Tsukuba, 305-8569, JAPAN

A WHOLE ORGANISM APPROACH TO THE PHYSIOLOGICAL IMPACTS OF OCEAN ACIDIFICATION. M53

Wood, Hannah L.¹, Stephen Widdicombe¹, and John I. Spicer²

Investigations into the impact of ocean acidification on an organism can be achieved by studying physiological processes. Currently, calcification and growth are popular proxies to quantify a species response because of their vulnerability to the changing carbonate chemistry predicted in a high CO₂ world. However, hypercapnia and acidosis have the potential to affect many other physiological processes within an organism. Moreover, these processes are frequently interdependent and by focusing on just one area (e.g. growth/calcification) there is a high possibility of overlooking sublethal impacts of ocean acidification. A series of mesocosm experiments have been conducted to investigate how key survival processes are affected by ocean acidification. These experiments have been conducted using the ophiuroid brittlestar *Amphiura filiformis*, a calcifying species which is an ecosystem engineer widespread across European temperate waters. Investigating how pH affects metabolism, ecosystem function (nutrient flux), reproduction, muscle maintenance, arm regeneration and differentiation, in addition to growth and calcification, gives a clear picture of how this organism is affected by, and potentially copes with ocean acidification. The results for *A. filiformis* show metabolic up-regulation and increased calcification, which result in a trade off with other physiological processes. While the results of the study indicate an increase in growth and

calcification, and therefore suggest no detrimental effect of increase pH, the whole organism approach reveals a different picture; the trade offs and prioritisation reveal the survival of this species in a high CO_2 ocean could be at risk.

RESPONSES OF PHYTOPLANKTON ASSEMBLAGE AND ORGANC CARBON DYNAMICS TO CO_2 INCREASE T53

<u>Yoshimura, Takeshi</u>¹, Jun Nishioka², Koji Suzuki³, Hiroshi Hattori⁴, Hiroshi Kiyosawa⁵, Daisuke Tsumune¹, Kazuhiro Misumi¹ and Takeshi Nakatsuka²

To investigate the responses of phytoplankton and organic carbon dynamics to CO₂ increase, a CO₂ manipulation experiment was conducted in the Okhotsk Sea in summer 2006. Surface water with a natural phytoplankton assemblage was incubated with bubbling air containing different CO₂ concentrations (180, 380, 750, and 1000 ppm). Temporal changes in phytoplankton pigments and particulate and dissolved organic carbon (POC and DOC) were observed for 14 days. The surface water was depleted in nutrients, so phytoplankton in the bottles remained at a low biomass under a regenerated system. If the values at the end of the experiment were compared in each treatment, the fucoxanthin/Chl-a ratios decreased with increasing CO₂, indicating the relative abundance of fucoxanthin-containing phytoplankton such as diatoms would be sensitive to a change in CO₂. Since no cells of coccolithophores were detected using a scanning electron microscope, we could not determine the response of them to the CO₂ gradient. The amount of DOC accumulation decreased with increasing CO₂, while no significant difference was observed for change in POC between treatments. The increase in atmospheric CO₂ is concluded to potentially affect the phytoplankton community structure and organic carbon flow in the nutrient-depleted subpolar surface water.

¹ Plymouth Marine Laboratory, Prospect Place, West Hoe, Plymouth PL1 3DH U.K.

² University of Plymouth, Drake Circus, Plymouth PL4 8AA hawo@pml.ac.uk

¹Central Research Institute of Electric Power Industry, 1646 Abiko, Abiko, Chiba 270-1194, JPN. (ytakeshi@criepi.denken.or.jp)

²Hokkaido University, North 19 West 8, Kita-ku, Sapporo, Hokkaido 060-0819, JPN.

³Hokkaido University, North 10 West 5, Kita-ku, Sapporo, Hokkaido 060-0810, JPN.

⁴Tokai University, 5-1-1-1 Minamisawa, Minami-ku, Sapporo, Hokkaido 005-8601, JPN.

⁵Marine Biological Research Institute of Japan, Shinagawa, Tokyo 142-0042, JPN.

Second Symposium on the Ocean in a High-CO₂ World List of Participants

Achterberg, Eric National Oceanography Centre University of Southampton European Way Southampton SO14 3ZH, U.K. eric@noc.soton.ac.uk

Alin, Simone NOAA-PMEL 7600 Sand Point Way NE Seattle, WA 98115, U.S.A. simone.r.alin@noaa.gov

Allemand, Denis Centre Scientifique de Monaco Avenue Saint-Martin MC-98000 Monaco allemand@centrescientifique.mc

Andersson, Andreas J Bermuda Institute of Ocean Sciences 17 Biological Station St. George's GE01, Bermuda andreas.andersson@bios.edu

Andersson, Pia M SMHI Sven Källfelts gata 15 42671 Gothenburg, Sweden pia.andersson@smhi.se

Anthony, Kenneth R The University of Queensland Centre for Marine Studies Gehrman Building 60 St Lucia, QLD 4072, Australia k.anthony@uq.edu.au

Appelhans, Yasmin S IFM-GEOMAR Düsternbrooker Weg 20 24105 Kiel, Germany yappelhans@ifm-geomar.de

Ardelan, Murat V Norwegian Univ. of Science and Technolgy Høgskolevein 5 7491 Trondheim, Norway murato@nt.ntnu.no Avgoustidi, Valia
Hellenic Center for Marine Research
47th km Athens - Sounio,
Mavro Lithari, PO Box 712
19013 Anavissos, Greece
vavgoustidi@ath.hcmr.gr

Azetsu-Scott, Kumiko
Bedford Institute of Oceanography
P.O. Box 1006
Dartmouth, NS B2Y 4A2, Canada
Azetsu-ScottK@mar.dfo-mpo.gc.ca

Barcelos e Ramos, Joana IFM-GEOMAR Düsternbrooker Weg 20 24105 Kiel, Germany jramos@ifm-geomar.de

Barry, James P MBARI 7700 Sandholdt Road Moss Landing, CA 95039, U.S.A. barry@mbari.org

Baxter, John Scottish Natural Heritage Silvan House 231 Corstorphine Road Edinburgh, EH12 7AT, U.K. John.Baxter@snh.gov.uk

Bednarsek, Nina British Antarctic Survey Madingley Road, High Cross Cambridge CB30ET, U.K. nindna@bas.ac.uk

Bernard, Olivier O INRIA, COMORE BP93 6902 Sophia-Antipolis Cedex, France olivier.bernard@inria.fr

Betti, Maria Marine Environment Laboratories Radiometrics Quai Antoine 1er MC-98000 Monaco m.betti@iaea.org Bijma, Jelle Alfred-Wegener-Institute Marine Biogeosciences Am Handelshafen 12 27570 Bremerhaven, Germany Jelle.Bijma@awi.de

Blackford, Jerry Plymouth Marine Laboratory Prospect Place Plymouth, PL1 3DH, U.K. jcb@pml.ac.uk

Blanchy, Bruno B
Gouvernement de Monaco
Département Equipement Environnement et
Urbanisme
10 bld de Belgique
MC 98000 MONACO
bblanchy@gouv.mc

Bleich, Markus Christian-Albrechts-University Kiel Department of Physiology Olshausenstr. 40 D-24098 Kiel, Germany m.bleich@physiologie.uni-kiel.de

Bock, Christian Alfred-Wegener-Institute Am Handelshafen 12 D-27570 Bremerhaven, Germany Christian.Bock@awi.de

Boetius, Antje MPI for Marine Microbiology Jacobs University Celsiusstr. 1 28359 Bremen, Germany aboetius@mpi-bremen.de

Boisson, Florence F International Atomic Energy Agency Marine Environment Laboratories 4 quai Antoine Ier MC 98000 Monaco F.Boisson@iaea.org

Boisson, Michel P.R. Scientific Center of Monaco "Les Villas des Pins" Bloc C 7, rue Honoré Labande MC 98000 Monaco mboisson@gouv.mc Bopp, Laurent
IPSL / LSCE
CE Saclay - Orme des Merisiers - Bat 712
F-91191 Gif sur Yvette, France
Laurent.Bopp@lsce.ipsl.fr

Borghi, Andrea Université de Neuchâtel Hydrogelogy Sur le Mont 32 1724 Praroman, Switzerland andrea.borghi@unine.ch

Børsheim, Knut Yngve Institute of Marine Research Group of Oceanography Nordnesgt 33 NO-5817 Bergen, Norway yngve.borsheim@imr.no

Bozec, Yann Station Biologique de Roscoff UMR 7144-Chimie Marine Place Georges Teissier BP74 29682 Roscoff, France bozec@sb-roscoff.fr

Breitbarth, Eike Gothenburg University Department of Chemistry Kemivaegen 10 SE-41296Gothenburg, Sweden eike@chem.gu.se

Breviere, Emily H SOLAS Project Office School of Environmental Sciences University of East Anglia Norwich NR4 7TJ, U.K. e.breviere@uea.ac.uk

Broadgate, Wendy J IGBP Secretariat Royal Swedish Academy of Sciences Box 50005 SE 104 05 tockholm, Sweden wendy@igbp.kva.se

Browning, David G University of Rhode Island Physics Department 2 Lippitt Road Kingston, RI 02881, U.S.A. decibeldb@aol.com Buxton, Lucy University of Technology Department of Environmental Science PO Box 123 Broadway Ultimo, Sydney, NSW 2007, Australia Lucy.Buxton@uts.edu.au

Caldeira, Ken Carnegie Institution 260 Panama St Stanford, CA 94305, U.S.A. kcaldeira@stanford.edu

Cassar, Nicolas Dept. of Geosciences, Guyot Hall Princeton University Princeton, New Jersey 08544, U.S.A. ncassar@princeton.edu

Catarino, Ana I Université Libre de Bruxelles Laboratoire de Biologie Marine ULB - Campus du Solbosch, CP 160/15 avenue F.D. Roosevelt 50 1050 Bruxelles, Belgium ana.catarino@ulb.ac.be

Chahine, Fanny
IAEA
Marine Environment Laboratories - RML
4 quai Antoine 1er
MC 98000 Monaco
F.Chahine@aiea.org

Charalampopoulou, Anastasia National Oceanography Centre European Way Southampton SO14 3ZH, U.K. ac4w07@soton.ac.uk

Chen, Baixin Heriot-Watt University School of Engineering and Physical Science Edinburgh EH14 5AS, U.K. B.Chen@hw.ac.uk

Chierici, Melissa University of Gothenburg Chemistry Department Kemivaegen 10 41296 Goteborg, Sweden melissa@chem.gu.se Chou, Lei
Université Libre de Bruxelles
Laboratoire d'Océanographie Chimique et
Géochimie des Eaux
ULB - CP 208, Campus de la Plaine
Boulveard du Triomphe
B-1050 Brussels, Belgium
lei.chou@ulb.ac.be

Clemmesen, Catriona Leibniz Institute for Marine Sciences IFM-GEOMAR Duesternbrooker Weg 20 24146 Kiel, Germany cclemmesen@ifm-geomar.de

Comeau, Steeve Laboratoire d'océanographie de Villefranche Chemin du Lazaret, station zoologique 6230 Villefranche sur Mer, France comeau@obs-vlfr.fr

Currie, Kim NIWA Centre for Chemical and Physical Oceanogr. Chemistry Dept, University of Otago PO Box 56 Dunedin 9054, New Zealand k.currie@niwa.co.nz

Cuschnir, Ariel A
The Louis Berger Group
2445 M Street
Washington DC, 20037, U.S.A.
acuschnir@louisberger.com

Czerny, Jan IFM-Geomar Biogeochemistry Duesternbroker Weg 20 24105 Kiel, Germany jczerny@ifm-geomar.de

Davenet, Valérie Direction de l'Environnement 3 avenue de Fontvieille MC 98000, Monaco vdavenet@gouv.mc

De Baar, Hein J Royal Netherlands Institute for Sea Research PO Box 59 1790 AB Den Burg, Texel, The Netherlands debaar@nioz.nl De Carlo, Eric Heinen University of Hawaii at Manoa Department of Oceanography Honolulu, HI 96822, U.S.A. edecarlo@soest.hawaii.edu

DeGrandpre, Michael D University of Montana Department of Chemistry 32 Campus Drive Missoula, MT 59812, U.S.A. michael.degrandpre@umontana.edu

Dickson, Andrew G Scripps Institution of Oceanograpy 9500 Gilman Drive La Jolla, California 92093, U.S.A. adickson@ucsd.edu

Dissard, Delphine Alfred Wegener Institute Am Handelshafen 12 Building E-2405 D-27520 Bremerhaven, Germany delphine.dissard@awi.de

Dubois, Philippe Université Libre de Bruxelles Biologie marine CP 160/15 ULB av Roosevelt 50 B-1050 Bruxelles, Belgium phdubois@ulb.ac.be

Dumousseaud, Cynthia C National Oceanography Centre European Way, Waterfront Campus Southampton SO14 3ZH, U.K. cd6@noc.soton.ac.uk

Dupont, Sam The Sven Lovén Centre for Marine Sciences 45034 Fiskebackskil, Sweden sam.dupont@marecol.gu.se

Egilsdottir, Hronn H University of Plymouth 7b Queens Gate Lipson, Plymouth PL4 7PW, U.K. hronne@gmail.com Emeka, Ozokolo I Lagos State University Civil Engineering 10 Alves street Lawanson, Surulere Lagos 23401, Nigeria ozoide2000@yahoo.co.uk

Enstad, Lars Inge L University of Bergen Allegaten 70 5007 Bergen, Norway lars.inge.enstad@bccs.uib.no

Erez, Jonathan The Hebrew University of Jerusalem Institute of Earth Sciences Givat Ram, Jerusalem, Israel 91904 erez@vms.huji.ac.il

Fabres, Joan J GRID-Arendal Marine Unit Teaterplassen 3 Arendal, Aust Agder 4836, Norway Joan.Fabres@grida.no

Fabry, Victoria J California State University San Marcos Department of Biological Sciences San Marcos, CA 92096, U.S.A. fabry@csusm.edu

Feely, Richard PMEL/NOAA - OCRD 7600 Sand Point Way NE Seattle, WA 98115, U.S.A. Richard.A.Feely@noaa.gov

Findlay, Helen S Plymouth Marine Laboratory Prospect Place, West Hoe Plymouth PL1 3DH, U.K. hefi@pml.ac.uk

Fine, Maoz Bar Ilan University Interuniversity Institute for Marine Science Eilat, 88103, Israel finema@mail.biu.ac.il Fiorini, Sarah S LOV 3, Rue de la Victoire 6230 Nice, France sarah.fiorini@obs-vlfr.fr

Fosså, Jan Helge J Institute of Marine Research P.O. Box 1870 Nordnes N-5817 Bergen, Norway jhf@imr.no

Fransson, Agneta University of Gothenburg Earth Science Centre, Oceanography Box 460 405 30 Goteborg, Sweden agneta@gvc.gu.se

Frommel, Andrea Y IfM-GEOMAR Duesternbrookerweg. 20 24105 Kiel, Germany afrommel@ifm-geomar.de

Gangstø, Reidun University of Bern Physics Institute, Sidlerstrasse 5 3012 Bern, Switzerland gangsto@climate.unibe.ch

Gattuso, Jean-Pierre CNRS, Laboratoire d'Océanographie BP 28 06234 Villefranche-sur-mer Cedex, France gattuso@obs-vlfr.fr

Gazeau, Frederic Nethelands Institute of Ecology P.O. Box 140 Yerseke, Zeeland 4400AC Netherlands f.gazeau@nioo.knaw.nl

Gehlen, Marion LSCE CEN Saclay - L'Orme des Merisiers Bât. 712 91191 Gif sur Yvette, France marion.gehlen@lsce.ipsl.fr Gorsky, Gabriel g CNRS / LOV B.P. 28 06234 Villefranche sur mer, France gorsky@obs-vlfr.fr

Goyet, Catherine UPVD 52 avenue Paul ALDUY 66860 Perpignan, France cgoyet@univ-perp.fr

Green, Alison J CSIRO PUBLISHING 150 Oxford Street Collingwood, 3066, Australia alison.green@csiro.au

Grosjean, Philippe P Universite de Mons-Hainaut Pentagone (3D08) 8, avenue du Champ de Mars Mons, Hainaut 7000, Belgium Philippe.Grosjean@umh.ac.be

Gross, Elizabeth SCOR College of Marine and Earth Studies Robinson Hall, University of Delaware Newark, DE 19716, U.S.A. egross@scor-int.org

Gruber, Nicolas
ETH Zurich
Department of Environmental Sciences
Universitätstrasse 16, CHN E 21.1
CH-8092 Zurich, Switzerland
nicolas.gruber@env.ethz.ch

Gutowska, Magdalena A Alfred-Wegener-Institute Marine Animal Physiology Bülow Str 7 24105 Kiel, Germany magdalena.gutowska@awi.de

Hall-Spencer, Jason M Marine Institute University of Plymouth Plymouth PL4 8AA, U.K. jhall-spencer@plymouth.ac.uk Hammer, Karen
Norwegian University of Science and
Technology
Biology Department
Brattørkaia 17b
7010 Trondheim, Norway
karenmh@stud.ntnu.no

Hansson, Lina Laboratoire d'Océanographie de Villefranche BP 28 06 234 Villefranche-sur-Mer, France hansson@obs-vlfr.fr

Harada, Koh
National Inst. of Advanced Industrial Science
and Technology
Research Institute for Environmental
Management Technology
16-1 Onogawa
Tsukuba, Ibaraki 305-8569, Japan
koh.harada@ni.aist.go.jp

Hauck, Judith Alfred-Wegener-Institute Marine Biogeosciences Am Schützenplatz 52 26121 Oldenburg, Germany judith.hauck@awi.de

Haugan, Peter M University of Bergen Geophysical Institute Allegaten 70 N-5007 Bergen, Norway Peter.Haugan@gfi.uib.no

Hauton, Chris C Unversity of Southampton National Ocanography Centre European Way Southampton SO14 3ZH, U.K. ch10@noc.soton.ac.uk

Havenhand, Jon N University of Gothenburg Tjärnö Marine Biological Laboratory 45296 Strömstad, Sweden jon.havenhand@marecol.gu.se Heilmayer, Olaf National Oceanography Centre, Southampton School of Ocean and Earth Science European Way Southampton, SO14 3ZH, U.K. oheilmayer@noc.soton.ac.uk

Heinrich, Hartmut
Federal Maritime and Hydrographic Agency
Marine Physics
Bernhard-Nocht-Strasse 78
20359 Hamburg, Germany
hartmut.heinrich@bsh.de

Heinze, Christoph University of Bergen Geophysical Institute and Bjerknes Centre Allégaten 70 5007 Bergen, Norway christoph.heinze@gfi.uib.no

Held, Hermann Potsdam Institute for Climate Impact Research PO Box 601203 14412 Potsdam, Germany held@pik-potsdam.de

Hood, Maria
UNESCO
Intergovernmental Oceanographic
Commission
1 Rue Miollis
Paris, 75732, France
m.hood@unesco.org

Hopkins, Frances E University of East Anglia School of Environmental Sciences Norwich NR4 7TJ, U.K. United Kingdom f.hopkins@uea.ac.uk

Houlbreque, Fanny Marine Environment Laboratory - IAEA 4 quai Antoine 1er BP 800 MC-98012 Monaco F.Houlbreque@iaea.org

Howard, William R Antarctic Climate & Ecosystems CRC Private Bag 80 Hobart, TAS 7001, Australia Will.Howard@acecrc.org.au Hu, Marian IfM-GEOMAR Biological Oceanography Hohenbergstr. 2 24105 Kiel, Germany mhu@ifm-geomar.de

Hydes, David J National Oceanography Centre Empress Dock Southampton SO14 3ZH, U.K. djh@noc.soton.ac.uk

Ianson, Debby
Institute of Ocean Sciences
9860 West Saanich Rd.
North Saanich, BC V8L 4B2, Canada
iansond@pac.dfo-mpo.gc.ca

Iglesias-Rodriguez, M. Debora D School of Ocean and Earth Science National Oceanography Centre European Way Southampton SO14 3ZH, U.K. dir@noc.soton.ac.uk

Ilyina, Tatjana University of Hawaii Department of Oceanography 1000 Pope Road Honolulu, HI 96822, U.S.A. ilyina@soest.hawaii.edu

Isensee, Kirsten
Leibniz Institute for Baltic Sea Research
Biological Oceanography
Seestrasse 15
18119 Rostock, Germany
kirsten.isensee@io-warnemuende.de

Ishida, Akio A Frontier Research Center for Global Change JAMSTEC 3173-25 Showamachi Kanazawa-ku, Yokohama, 236-0001, Japan ishidaa@jamstec.go.jp

Ishida, Hiroshi H
The General Environmental Technos CO.,
1-3-5, Azuchimachi, Chuo-ku
Osaka, 541-0052, Japan
ishida hiroshi@kanso.co.jp

Ishii, Masao Meteorological Research Institute Geochemical Research Department 1-1 Nagamine Tsukuba, Ibaraki 305-0052, Japan mishii@mri-jma.go.jp

Jeffree, Ross A
IAEA
Marine Environment Laboratories
4 Quai Antoine 1er
MC 98000 Monaco
R.Jeffree@iaea.org

Kasemann, Simone A University of Edinburgh School of Geosciences West Mains Road Edinburgh EH16 5NF, U.K. simone.kasemann@ed.ac.uk

Keeling, Ralph Scripps Institution of Oceanography 9500 Gilman Drive, #0210 La Jolla, CA 92093, U.S.A. rkeeling@ucsd.edu

Kellermann, Adolf ICES H.C. Andersens Boulevard 44-46 DK-1553 Copenhagen, Denmark adi@ices.dk

Kerrison, Philip D University of Essex Biological Science Wivenhoe Park Colchester CO4 3SQ, U.K. pkerri@essex.ac.uk

Kleypas, Joanie
National Center for Atmospheric Research
Institute for the Study of Society and
Environment
PO Box 3000
Boulder, CO 80307, U.S.A.
kleypas@ucar.edu

Knoll, Andrew H Harvard University Organismic and Evolutionary Biology 26 Oxford Street Cambridge, MA 02138, U.S.A. aknoll@oeb.harvard.edu Kosakowska, Alicja
Institute of Oceanology
Marine Chemistry and Biochemistry
Department
Powstancow Warszawy 55
81-712 SOPOT, Poland
akosak@iopan.gda.pl

Kristiansen, Erlend NTNU Department of Biology Høgskoleringen 5 7491 Trondheim, Norway Erlend.Kristiansen@bio.ntnu.no

Krug, Sebastian A IFM-GEOMAR RD2 - Marine Biogeochemistry Düsternbrooker Weg 20 24105 Kiel, Germany skrug@ifm-geomar.de

Kulkarni, Balasaheb G
The Institute of Science
Zoology, Environmental science
301 Vinbhavri, Dinsha vachha Road
Churchgate, Mumbai 400 020
India
balasahebk@yahoo.com

Kurihara, Haruko Nagasaki University Institute for East China Sea Research Tairamachi 1551-7 Nagasaki, Nagasaki 851-2213, Japan harukoku@e-mail.jp

Lacoue-Labarthe, Thomas Radioecology Laboratory REL-MEL, IAEA 4 Quai Antoine Ier MC 98 000 MONACO, tlacouel@univ-lr.fr

Langdon, Chis University of Miami Marine Biology and Fisheries 4600 Rickenbacker Cswy Miami, FL 33149, U.S.A. clangdon@rsmas.miami.edu Laverock, Bonnie B Plymouth Marine Laboratory Prospect Place, West Hoe Plymouth PL4 6NN, U.K. bonver@pml.ac.uk

Leblud, Julien Université Mons-Hainaut Numerical Ecology of Aquatic Systems 128, rue A.Descamps 7021 HAVRE, Belgium julien.leblud@umh.ac.be

Lehodey, Patrick CLS Satellite Oceanography Division 8-10 rue Hermes 31520 Ramonville, France plehodey@cls.fr

Leinen, Margaret Climos, Inc. 119 S Columbus Street Alexandria, VA 22314, U.S.A. mleinen@climos.com

Levitan, Orly
Bar Ilan University
The Mina and Everard Goddman Faculty of
Life Sciences
31/2 Pines St.
Neve Zedek, Tel Aviv, 65135, Israel
levitao@mail.biu.ac.il

Lianou, Vasiliki University of Athens Geology Megaloupoleos 43 11476 Athens, Greece vlianou@geol.uoa.gr

Lipschultz, Fred NASA Headquarters 300 E St. SW Washington D.C., 20546, U.S.A. fredric.lipschultz@nasa.gov

Liu, Jinwen CNRS-LOV Laboratoire d'Océanographie BP28 06234 Villefranche-sur-Mer, France liu@obs-vlfr.fr MacCracken, Michael C Climate Institute 6308 Berkshire Drive Bethesda, MD 20814, U.S.A. mmaccrac@comcast.net

Mahanta, Chandan Indian Institute of Technology Guwahati Civil Engineering Guwahati, Assam 781039, India mahanta iit@yahoo.com

Maier, Cornelia Laboratoire d'Océanographie de Villefranche Microbial Ecology and Biogeochemistry Group BP 28

06234 Villefranche-sur-mer, France maier@obs-vlfr.fr

Marinelli, Roberta L National Science Foundation Antarctic Sciences Division 4201 Wilson Blvd Arlington, VA 22230, U.S.A. rmarinel@nsf.gov

Martin, Sophie International Atomic Energy Agency Marine Environment Laboratories BP 800 MC-98012 Monaco S.Martin@iaea.org

Marubini, Francesca
Joint Nature Conservation Committee
Dunnet House
7, Thistle Place
Aberdeen AB10 1UZ, U.K.
francesca.marubini@jncc.gov.uk

Masuda, Yoshio Hokkaido Univ. N10 W5, Kita-ku Sapporo, Hokkaido 060-0810, Japan y-masuda@ees.hokudai.ac.jp

Matear, Richard J CSIRO GPO Box 1538 Hobart, TAS 7001, Australia richard.matear@csiro.au Mathis, Jeremy T University of Alaska Fairbanks Institute of Marine Science 245 O'Neil Fairbanks, AK 99775, U.S.A. jmathis@sfos.uaf.edu

Maxon, Mary Gordon and Betty Moore Foundation The Presidio of San Francisco PO Box 29910 San Francisco, CA 94129, USA mary.maxon@moore.org

McGrath, Triona T Marine Institute Rinville Oranmore, Co.Galway, Ireland triona.mcgrath@marine.ie

McGovern, Evin E Marine Institute Rinville Oranmore, Co.Galway, Ireland evin.mcgovern@marine.ie

McGraw, Christina University of Otago Department of Chemistry Dunedin, New Zealand cmcgraw@chemistry.otago.ac.nz

McKinnell, Skip PICES / IOS 9860 W. Saanich Rd. North Saanich, BC, BC V8L 4B2, Canada mckinnell@pices.int

McNeil, Ben Climate Change Research Centre University of New South Wales Sydney, NSW 2040, Australia b.mcneil@unsw.edu.au

Melzner, Frank IFM-GEOMAR Biological Oceanography Hohenberg Str. 2 24105 Kiel, Germany fmelzner@ifm-geomar.de Merico, Agostino A GKSS Research Centre Institute for Coastal Research Max Planck Strasse 1 21502 Geesthacht, Germany agostino.merico@gkss.de

Miest, Joanna J Sierichstrasse 102 22299 Hamburg, Germany jo.miest@web.de

Miquel, Juan Carlos IAEA Marine Environment Laboratories 4 Quai Antoine 1er MC-98000 Monaco j.c.miquel@iaea.org

Monteiro, Pedro P CSIR / Oceans and Climate Jan Cilliers St Stellenbosch 7599,South Africa pmonteir@csir.co.za

François Morel Princeton University Guyot Hall Princeton, NJ, 08544 U.S.A. morel@princeton.edu

Moya, Aurélie EA 4228 ECOMERS, University of Nice Sophia Antipolis Avenue Valrose, BP 71 6108 Nice cedex 2, France aurelie.moya@unice.fr

Müller, Marius N IfM-Geomar Marine Biogeochemistry Düsternbrooker Weg 20 24105 Kiel, Germany mnmueller@ifm-geomar.de

Munhoven, Guy University of Liège Inst. d'Astrophysique et de Géophysique 17 Allée du Six-Août 4000 Liège, Belgium Guy.Munhoven@ulg.ac.be Akihiko Murata JAMSTEC 2-15, Natsushima-cho Yokosuka Kanagawa 237-0061, Japan murataa@jamstec.go.jp +81-468-67-9503

Murray, James W University of Washington School of Oceanography Box 355351 Seattle, WA 98195, U.S.A. jmurray@u.washington.edu

Nisumaa, Anne-Marin A LOV-Laboratoire d'Océanographie BP 28 06234 Villefranche sur Mer, France nisumaa@obs-vlfr.fr

Nojiri, Yukihiro National Institute for Environmental Studies Center for Global Environment Research 16-2, Onogawa Tsukuba, Ibaraki 305-8506, Japan nojiri@nies.go.jp

Oechel, Walter C San Diego State University Biology/Ecology 5500 Campanile Drive San Diego, CA 92182, U.S.A. oechel@sunstroke@sdsu.edu

Olafsdottir, Solveig Marine Research Institute Skulagata 4 IS 105 Reykjavi, Iceland solveig@hafro.is

Olafsson, Jon University of Iceland Marine Research Institute Skulagata 4 IS 121 Reykjavik, Iceland jon@hafro.is

Omstedt, Anders A University of Gothenburg Earth Sciences: Oceanography Box 460 SE-405 30 Gothenburg, Sweden Anders.Omstedt@gvc.gu.se Orlikowska, Anna
Leibniz Institute for Baltic Sea Research
Warnemünde (IOW)
Department of Marine Chemistry
Seestraße 15
18119 Rostock-Warnemünde, Germany
anna.orlikowska@io-warnemuende.de

Orr, James C IAEA, Marine Environmental Labs 4 Quai Antoine 1er MC-98000 Monaco j.orr@iaea.org

Ronald Osinga Wageningen University PO Box 338 6700 AH Wageningen, Netherlands ronald.osinga@wur.nl

Ounais, Nadia Musée Océanographique Avenue Saint-Martin MC-98000 Monaco n.ounais@oceano.mc

Pansch, Christian IFM-GEOMAR Marine Ecology - Experimental Ecology Düsternbrooker Weg 20 24105 Kiel, Germany cpansch@ifm-geomar.de

Pantoja, Silvio University of Concepcion Oceanografia, Barrio Universitario Concepcion, Chile spantoja@udec.cl

Park, Susan National Academy of Sciences Ocean Studies Board 500 5th Street, NW Washington, DC 20001, U.S.A. spark@nas.edu

Parker, Laura University of Western Sydney School of Natural Sciences Locked Bag 1797, Penrith South DC Sydney, NSW 1797, Australia lauraparker 83@hotmail.com Pedersen, Sindre Andre NTNU, Biology Høgskoleringen 7491 Trondheim, Norway sindre.pedersen@bio.ntnu.no

Perttila, Matti M Finnish Institute of Marine Research Erik Palmenin aukio 1 FIN-00561 Helsinki, Finland matti.perttila@fimr.fi

Piontek, Judith Alfred-Wegener-Institute Am Handelshafen 12 27570 Bremerhaven, Germany Judith.Piontek@awi.de

Plattner, Gian-Kasper ETH Zurich Department of Environmental Sciences Universitätstrasse 16, CH-8092 Zurich, Switzerland gian-kasper.plattner@env.ethz.ch

Poertner, Hans O Alfred-Wegener-Institute Marine Animal Physiology Am Handelhafen 12 27570 Bremerhaven, Germany hans.poertner@awi.de

Porzio, Lucia Stazione Zoologica "A. Dohrn" Via Sant'Abbondio 132 80045 Pompei, Italy lucia.porzio@szn.it

Premovic, Pavle I University of Nis Laboratory for Geochemistry P. O. Box 224, 18000 Nis, Yugoslavia asteroid.pavle@yahoo.com

Prol-Ledesma, Rosa Maria Universidad Nacional Autonoma de Mexico Instituto de Geofísica Cd. Universitaria, Coyoacan Mexico City, D.F. 4510, Mexico Prol@geofisica.unam.mx Ransome, Emma J Plymouth Marine Laboratory Microbiology 13b Alexandra Road Mutley, Plymouth PL4 7EE, U.K. emmajane7@hotmail.com

Reynaud, Stéphanie Centre Scientifique de Monaco Av Saint Martin MC-8000 MONACO sreynaud@centrescientifique.mc

Rice, Donald L National Science Foundation Division of Ocean Sciences 4201 Wilson Boulevard Arlington, VA 22230, U.S.A. drice@nsf.gov

Richier, Sophie UMR7093 CNRS 181, chemin du Lazaret 06234 Villefranche-sur-Mer, France richier@obs-vlfr.fr

Riddle, Martin Australian Antarctic Division Environmental Protection and Change 203 Channel Highway Kingston, TAS 7050, Australia martin.riddle@aad.gov.au

Ridgwell, Andrew J University of Bristol School of Geographical Sciences University Road Bristol BS8 1SS, U.K. andy@seao2.org

Riebesell, Ulf Leibniz Institute of Marine Sciences IFM-GEOMAR Biological Oceanography Duesternbrooker Weg 20 24105 Kiel, Germany uriebesell@ifm-geomar.de

Robbins, Lisa L US Geological Survey 600 4th St South St Petersburg, FL 33701, U.S.A. lrobbins@usgs.gov Roberts, Donna Antarctic Climate & Ecosystem CRC Private Bag 80 Hobart, TAS 7001, Australia d.roberts@acecrc.org.au

Rodolfo-Metalpa, Riccardo University of Plymouth, Marine Institute Plymouth, PL4 8AA, U.K. riccardo.rodolfo-metalpa@plymouth.ac.uk

Russell, Joellen L University of Arizona Department of Geosciences 1040 E 4th St Tucson, AZ 85721, U.S.A. jrussell@email.arizona.edu

Sabine, Christopher L NOAA/PMEL 7600 Sand Point Way NE Seattle, Washington 98115, U.S.A. chris.sabine@noaa.gov

Sanchez Cabeza, Joan-Albert IAEA, Marine Environment Laboratories 4, Quai Antoine 1er MC-98000 MONACO j.a.sanchez@iaea.org

Saphörster, Julia IFM-GEOMAR Hohenbergstr. 2 24105 Kiel, Germany jsaphoerster@ifm-geomar.de

Sasai, Yoshikazu
Japan Agency for Marine-Earth Science and
Technology
3173-25, Showa-machi, Kanazawa-ku,
Yokohama, 236-0001, Japan
ysasai@jamstec.go.jp

Sato, Toru
University of Tokyo
Dept of Ocean Technology, Policy, and
Environment
Rm 474, Env. Build.
5-1-5 Kashiwa-n-ha
Kashiwa, 277-8563, Japan
sato-t@k.u-tokyo.ac.jp

Sawada, Yoshifumi S Kinki University Fisheries Laborotory Ohshima 1790-4 Kushimoto, Wakayama 649-3633, Japan YoshifumiSawada@za.ztv.ne.jp

Schmidt, Daniela N University of Bristol Earth Sciences Wills Memorial Building Queens Road Bristol, BS8 1RJ, U.K. d.schmidt@bristol.ac.uk.

Schneider, Birgit University of Kiel Ludewig-Meyn-Str. 10 24098 Kiel, Germany bschneider@gpi.uni-kiel.de

Schulz, Kai K IFM-GEOMAR Biogeosciences Düsternbrooker Weg 20 24105 Kiel, Germany kschulz@ifm-geomar.de

Schutter, Miriam Wageningen University Marijkeweg 40 6701 BM Wageningen, Netherlands miriam.schutter@wur.nl

Segschneider, Joachim Max-Planck-Institute for Meteorology Bundesstrasse 53 22605 Hamburg, Germany joachim.segschneider@zmaw.de

Shadwick, Elizabeth H. Dalhousie University Oceanography Department Halifax, NS B3H 4J1, Canada elizabeth.shadwick@dal.ca

Shin, Kyoungsoon Korea Ocean Research and Development Southern Coastal Environment Research Division 391 Jangmok-ri Jangmok-myon Geoje, 656-830 Soth Korea ksshin@kordi.re.kr Shirayama, Yoshihisa Kyoto University Field Science Education and Research Center 459 Shirahama, Wakayama 649-2211, Japan yshira@bigfoot.com

Shitashima, Kiminori
Central Research Institute of Electric Power
Industry
Environmental Science Research Laboratory
1646, Abiko
Abiko, Chiba 270-1194, Japan
shita@criepi.denken.or.jp

Simonet, Raphaël Direction de l'Environnement 3 avenue de Fontvieille MC-98000 Monaco rsimonet@gouv.mc

Smit, Tim Utrecht University Van der Horststraat 17 3141TH Maassluis, The Netherlands t.smit@students.uu.nl

Smythe-Wright, Denise National Oceanography Centre European Way Southampton, SO14 3ZH, U.K. dsw@noc.soton.ac.uk

Stumpp, Meike IFM-GEOMAR Biological Oceanography Hohenberg Str. 2 24105 Kiel, Germany mstumpp@ifm-geomar.de

Suatoni, Lisa Natural Resources Defense Council 40 W. 20th St. NY, NY 10011, U.S.A. Isuatoni@nrdc.org

Suckling, Coleen C British Antarctic Society Biological Sciences Dept. 8C Longsdale Terrace Oban, PA34 5JS, U.K. coleenclaire@yahoo.co.uk Sudhaus, Stefanie IFM-GEOMAR Leibniz Institute of Marine Sciences Düsternbrooker Weg 20 24105 Kiel, Germany ssudhaus@ifm-geomar.de

Suggett, David J Department of Biological Sciences University of Essex Colchester, CO4 3SQ, U.K. dsuggett@essex.ac.uk

Sundby, Bjorn Earth & Planetary Sciences McGill University Montreal, QC H3A 2A7, Canada bjorn.sundby@mcgill.ca

Suzuki, Toru Japan Hydrographic Association Marine Information Research Center Daiichi Sogo Bldg. 6F, 1-6-6, Hanedakuko Ota-ku, Tokyo 144-0041, Japan suzuki@mirc.jha.jp

Tambutté, Sylvie S Centre Scientifique de Monaco Physiology/Biochemistry Avenue Saint Martin MC-98000 Monaco stambutte@centrescientifique.mc

Taylor, Phillip R U.S. National Science Foundation Division of Ocean Sciences 4201 Wilson Blvd., Suite 725 Arlington, Virginia 22201, U.S.A. prtaylor@nsf.gov

Teyssie, Jean Louis IAEA-MEL 4 quai Antoine premier MC-98000 MONACO j.teyssie@iaea.org

Thatje, Sven National Oceanography Centre School of Ocean and Earth Science European Way Southampton, SO14 3ZH, U.K. svth@noc.soton.ac.uk Thorndyke, Michael C Sven Lovén Centre for Marine Scieces Kristineberg 450 34 Fiskebackskil, Sweden mike.thorndyke@marecol.gu.se

Thuesen, Erik V Evergreen State College 2700 Evergreen Parkway NW Lab I Olympia, WA 98505, U.S.A. thuesene@evergreen.edu

Touratier, Franck Université de Perpignan 52 av Paul Alduy 66860 Perpignan, France touratie@univ-perp.fr

Towanda, Trisha
Evergreen State College/University of Rhode
Island
1407 Kaiser Rd SW
Olympia, WA 98512, U.S.A.
trisha towanda@yahoo.com

Triantafyllou, George N Hellenic Centre for Marine Research Institute of Oceanography P.O.Box 712 Mavro Lithari Anavissos, Attica 19013, Greece gt@ath.hcmr.gr

Trull, Thomas W University of Tasmania, CSIRO ACE CRC PB 80 Hobart, TAS 7001, Australia Tom.Trull@utas.edu.au

Turley, Carol Plymouth Marine Laboratory Prospect Place, The Hoe Plymouth, PL1 3DH, U.K. ct@pml.ac.uk

Tyrrell, Toby
National Oceanography Centre
Southampton University
European Way
Southampton, SO14 3ZH, U.K.
tt@noc.soton.ac.uk

Urban, Ed

Scientific Committee on Oceanic Research College of Marine and Earth Studies Robinson Hall, University of Delaware Newark, DE 19716, U.S.A. Ed.Urban@scor-int.org

Vance, Thomas University of Newcastle upon Tyne Flat 1, 53 Percey Park Tynemouth, NE30 4JX, U.K. thomas.vance@newcastle.ac.uk

Veglia, André A
Direction de l'Environnement
Equipement, Environnement et Urbanisme
3, avenue de Fontvieille
MC-98000 Monaco
aveglia@gouv.mc

Wanninkhof, Rik NOAA/AOML 4301 Rickenbacker Causeway Miami, Florida 33149, U.S.A. rik.wanninkhof@noaa.gov

Watanabe, Yuji
The General Environmental Technos Co., Ltd.
Depart. Environement
1-3-5, Azuchimachi, Chuo-ku
Osaka, 541-0052, Japan
watanabe yuji@kanso.co.jp

Watson, Sue-Ann School of Ocean and Earth Sciences National Oceanography Centre European Way Southampton, SO14 3ZH, U.K. suwa@noc.soton.ac.uk

Weinbauer, Markus Laboratoire d'Océanographie BP 28 6234 0-sur-Mer, France wein@obs-vlfr.fr

Widdicombe, Stephen Plymouth Marine Laboratory Prospect Place, West Hoe Plymouth, Devon PL1 3DH, U.K. swi@pml.ac.uk Williams, Mary Ann
International Geosphere-Biosphere
Programme
Lilla Frescativägen 4
11418 Stockholm, Sweden
maryann@igbp.kva.se

Williams, Susan F UNESCO Bureau of Public Information 7 Place de Fontenoy 75005 Paris, France s.williams@unesco.org

Wood, Hannah L Plymouth Marine Laboratory Prospect Place West Hoe Plymouth, PL1 3DH, U.K. hawo@pml.ac.uk

Wood, Rachel University of Edinburgh Grant Institute West Mains Road Edinburgh, EH9 3JW, U.K. Rachel.Wood@ed.ac.uk

Yates, Kimberly K U.S. Geological Survey 600 Fourth Street South St. Petersburg, FL 33701, U.S.A. kyates@usgs.gov

Yoshimura, Takeshi Central Research Institute of Electric Power ndustry Environmental Science Research Laboratory 1646 Abiko Abiko, Chiba 270-1194, Japan ytakeshi@criepi.denken.or.jp

Zeebe, Richard E University of Hawaii School of Ocean and Earth Science and Technology 1000 Pope Rd, MSB 504 Honolulu, HI 96822, U.S.A. zeebe@hawaii.edu

Second Symposium on the Ocean in a High-CO2 World Author Index

Achterberg, Eric	44, 51, 59, 61	Buia, Maria Cristina	75
Albertson, Skip L.	37	Prol-Ledesma, Rosa Ma.	75
Alendal, Guttorm	52	Bustamante, Paco	65
Alin, Simone R.	37	Buxton, Lucy	43
Allemand, Denis	10, 24	Byrne, Robert H.	15
Anderson, Leif G.	73	Cabioch, Guy	50
Andersson, Andreas J.	12, 33, 48	Cadule, P.	54
Anthony, Ken	37, 71		11, 16, 20
Ardelan, Murat V.	23, 38	Campbell, Douglas	66
Atsushi, Ishimatsu	62	Canet, Carles	75
Auger, Hélène	51	Catarino, Ana I.	44, 55
Avgoustidi, V.	38, 86	Charalampopoulou, Anastasia	44
Aydin, Kerim	29	Chen, Baixin	45
Badger M.	43	Chou, Lei	45
Bakker, Dorothee	40, 59	Chourrout, Daniel	51
Barcelos e Ramos, Joana	39, 47	Christaki, U.	38
Barry, James P.	40	Christian, Jim	29
Bates, Nicholas R.	33, 68	Clemmesen, Catriona	41, 46, 59
Batigny, A.	65	Comeau, Steeve	46, 67
Baumert, Gunnar A.	41	Cornelia, Maier	67
Baxter, John	13, 35	Croot, Peter L.	23
Beaumont, Laurence	43	Cullison, Sarah	48
Bednarsek, Nina	40	Currie, Kim I.	47, 69
Bellerby, Richard J.	23, 28, 57, 70	Czerny, Jan T. B.	47
Benoit-Cattin, Alice	22	Dando, Paul R.	75
Bentov, Shmuel	24	Danguy, Théodore	43
Berman-Frank, Ilana	66	Danielsen, Magnus	22
Bernard, O.	11, 25	DaRocha, Regine	50
Berry, Katherine	77	Davies, Diana	87
Bijma, Jelle	50	Davis, Nancy	29
Blackford, Jerry	41, 59	De Baar, Hein	10, 23
Blain, Stéphane	43	de Beer, Dirk	26
Bleich, Markus	41	De Bodt, Caroline	45
Bock, Christian	49	De Carlo, Eric H	48
Boetius, Antje	11, 26	de Vargas, Colomban	76
Boisson, Florence	42, 67	DeGrandpre, Mike	48
	12, 33, 54, 73, 81	Deigweiher, Katrin	49
Borchard, Corinna	45, 74	Delille, Bruno	45
Borges, Alberto V.	45	Diaz-Pulido, Guillermo	37
Bouquet, Jean-Marie	51	Dickson, Andrew G.	49
Bozec, Yann	43	Dissard, Delphine	50
Braun, Alon	24	Douville, Eric	50
Bray, Stephen G.	20, 21	Dubois, Philippe	44, 55
Breitbarth, Eike	10, 23	Dumousseaud, Cynthia	44, 51
Brewer, Peter	45	Dupont, Sam	51, 58, 85
Brown, M. Christopher	66	Eakin, C. Mark	17
Browning, David	12, 32	Economou, S.	38
Bucciarelli, Eva	43	Edenhofer, Ottmar	35
Buck, Kurt R.	40	Egashira Takeshi	28
Budd, David A.	17	Eggins, Stephen	19
,	1,	200, 500 p.1011	1)

Eisenhauer, Anton	41, 45	Harlay, Jérôme	45
Engel, Anja	74	Hashioka, Taketo	56
Enstad, Lars Inge	52	Hattori, Hiroshi	89
Erez, Jonathan	10, 24	Hauck, Judith	57
Eric F. Pane	40	Haugan, Peter M.	10, 11, 30, 52
Fabry, Victoria	10, 11, 19, 73	Hauri Claudine	16, 11, 30, 32
Falkowski, Paul G.	76	Hauton, Chris	57
Fallick, Anthony E.	21	Havenhand, Jon	58, 85
	, 15, 37, 48, 78	Hayashi, Masahiro	62
Ferrier-Pagès, Christine	24	Hegeman, Jan	67
Ferrier-Pagès, Christine	76	Heilmayer, Olaf	85
Fielding, Sophie	40	Heinemann, Agnes	79
	52		58, 60
Findlay, H.S., Form, Armin	41	Heinze, Christoph Heip, Carlo	54
· · · · · · · · · · · · · · · · · · ·	11, 28	* '	
Fosså, Jan Helge Frank, Norbert	50	Held, Hermann Hepburn, Christopher D.	13, 35 69
Franke, Andrea	41, 46	Hepburn, Leanna	63
Frenzel, Hartmut	16 48	Hernandez-Ayon, J. Martin	78
Friederich, Gernot		Hiebenthal, Class	41
Frölicher, Thomas	73	Higuchi, Kazuhiro	80
Frommel, Andrea Y.	59 50	Hönisch, Bärbel	19
Gaillardet, Jérôme	50 53	Honryo, Tomoki	80
Gangstø, Reidun	53	Hopcroft, Russell R.	20
Gattuso, J-P. 9, 17, 46, 53		Hopkins, Frances E.	12, 34
Gazeau, Frédéric	54	Hoppema, Mario	57
Gehlen, Marion	53, 54, 81	Houlbreque, Fanny	67
Geider, Richard J.	84	Howard, William R.	10, 20, 21
Gerringa, Loes	23	Hu, Marian YA.,	59
Glynn, Peter W.	17	Hunter, Keith A.	47, 69
Goodman, Paul J.	78	Hurd, Catriona L.	69
Gorsky, Gaby	46	Hydes, David	51, 59
Goyet, C.	86	Hyun, Bong-Gil	83
Greeley, Dana	15	Ianson, Debby	78
Green, Darryl	44	Ilyina, Tatjana,	60
Grinstein, Mor	24	Inagaki, Fumio	26
Groom, Steve	45	Inoue, Hisayuki Y.	62
Grosjean, A. Ph.	65	Isensee, Kirsten	60
Grossteffan, Emilie	43	Ishida, Akio	61, 79
Gruber, Nicolas	9, 15, 16	Ishida, Hiroshi	61
Guibourt, Virginie	55	Ishii, Masao	62
Guillot, Antoine	43	Ishimatsu, Atsushi	64
Guillou, Jacques	43	Jakobsen, Tore	28
Gustafsson, Erik	72	Jeffree, Ross	42, 46, 65, 67
Gutowska, Magdalena A.	55, 70	Johansen, Henning	60
Haeckel, Matthias	26	Jokiel, Paul L.	33
Haga, Yutaka	80	Joly, Jean-Stéphane	51
Hales, Burke	78	Jones, Peter	73
Hallock, P.	77	Joos, Fortunat	15, 53, 73
Hall-Spencer, Jason M.	11, 29, 75	Juillet-Leclerc, Anne	50
Händel, Nicole	45, 74	Jutterström, Sara	73
Hanel, Reinhold	41	Kano, Yuki	11, 30
Hansson, L.	53	Kasemann, Simone A.	10, 21, 81
Harada, Koh	56, 88	Kendall, M.A.	52
Haramaty, Liti	76 51 50	Kerrison, Philip	63
Hardman-Mountford, Nick	51, 59	Kerros, Marie-Emmanuelle	76

Vimata Hidashi	28	Melzner, Frank 41, 55, 59, 70, 79, 84
Kimoto, Hideshi Kinoshita, Katsumoto	28	Melzner, Frank 41, 55, 59, 70, 79, 84 Mempel, Helgi 41
Kita, Jun	30	Merlivat, Liliane 43
Kita, Jun Kiyosawa, Hiroshi	89	Mestre, Nelia 85
Kleypas, Joan A.	9, 10, 17	Meyerhöfer, Michael 23
Kline, David	71	Michalopoulos, N. 38
Klinger, Terrie	72	Middelburg, Jack 54
Knoll, Andrew H.	12, 31	Midorikawa, Takashi 62
Knorr, P.	77	Millero, Frank J. 15
Knut Yngve Børsheim	43	Misumi, Kazuhiro 89
Kojima, Yuki	62	Möller, Volker 41
Konstantinopoulou, A.	38	Montagna, Paolo 76
Kosakowska, Alicja	63	Monteiro, P.M.S. 70
Krasakopoulou, E.	38, 86	Mooney, Ann 71
Krug, Sebastian A.	64	Moore, S. 82
Kuffner, Ilsa B.	33	Mowlem, Matt 51, 59
Kumiko, Azetsu-Scott	39	Moy, Andrew D. 10, 20, 21
Kurihara, Haruko	64	Moy, Andrew D., 20
Lachkar, Zouhair	16	Müller, Marius N. 12, 31, 39
Lacoue-Labarthe, Thomas	65, 67	Munhoven, Guy 71
Laffoley, Dan	35	Murray, James W. 72
Langdon, Chris	17	Nakadate, Akira 62
Langenbuch, Martina	55, 70	Nakamura, Koichi 26
LaRoche, Julie	66	Nakatsuka, Takeshi 89
Lawson, Tracy	84	Nausch, Monika 60
Lazar, Boaz	24	Nehrke, Gernot 50
Leblud, J.	65	Neill, Craig C. 23
Lee, Kitack	83	Newton, Jan 37
Lefebvre, Stephane	84	Nightingale, Philip D. 34
Lehodey, Patrick	11	Nishio, Masahiro 45
Leong, D.	82	Nishioka, Jun 89
Lerman, Abraham	33	Nojiri, Yukihiro 11, 28
Lombard, Fabien	19	Nzigou, Aimé Roger 54
Louvat, Pascal	50	O'Connor, Wayne A. 18
Lovera, Chris	40	Oberhansli, François 42, 65, 67
Lucas, Mike	44	Okada, Tokihiko 80
Lucassen, Magnus	49	Okunishi, Takeshi 56
Lunau, Mirko	74	Olafsdottir, Solveig R. 22
Mackas, David	29	Olafsson, Jon 10, 22
Mackenzie, Fred T.	33, 48	Omstedt, A, 72
Maeda, Nobuhiro	88	Orlikowska, Anna 73
Magi, Michimasa	61	Orly, Levitan 66
Mahanta, Chandan	66	Orr, James C. 9, 13, 15, 54, 67, 73
Maier-Reimer, Ernst	60, 73, 82	Osseforth, Christian 85
Manzello, Derek P.	17	Pantoja, Silvio 10
Martin, Sophie	65, 67, 76	Papathanassiou, E. 38
Marubini, Francesca,	10, 24	Parker, Laura M. 9, 18
Masuda, Yoshio	68	Paterne, Martine 50
Matear, Richard J.	68, 69	Paterson, Kristina 77
Mathis, Jeremy T.	68	Peacock, Cynthia G. 37
Matthiessen, Birte	64	Peck, Lloyd S. 19
McCulloch, Malcolm	76	Pempkowiak, Janusz 63
McGraw, Christina M.	69	Perttilä, Matti 74
McKinnell, Skip	11, 29	Peter S. Liss 34
McNeil, Ben I.	68, 69	Piatkowski, Uwe 41

Piepenburg, Dieter	41	Shadwick, E.H.	82
Piontek, Judith	45, 74	Shillito, Bruce	85
Plattner, Gian-Kasper	9, 15, 16	Shin, Kyoungsoon,	83
Pollani, A.	86	Shirayama, Yoshihisa	28, 61
Pörtner, Hans O.	11, 25, 49, 55	Shitashima, Kiminori	83
Porzio, Lucia	75	Silverman, Jack	24
Poulton, Alex	44	Slauenwhite, David	39
Prave, Anthony R.	21	Smit, Tim	87
Prowe, E.F.	82	Snape, Ian	77
Rabouille, S.	25	Sommer, Frank	41
Raines, Christine	84	Sommer, Ulirch	41, 64
Ralph, P.	43	Spero, Howard J.	19
-	69	<u>-</u>	
Raven, John A.		Spicer, John I.	27, 52, 88
Rehder, Gregor	26	Spindler, Michael	41
Reid, Malcolm	47	Steinacher, Marco	15, 73
Répécaud, Michel	43	Steinke, Michael	34, 63
Reynaud, Stéphanie	76	Steinnes, Eiliv	38
Richier, Sophie	76	Strassmann, Kuno M.	15
Riddle, Martin J.	77	Stumpp, Meike	84
Ridgwell, Andy	12, 34, 81	Sudhaus, Stefanie	66
Riebesell, Ulf 11, 12,	13, 23, 26, 31	Suggett, David J.	63, 84
32, 39.	41, 47, 60, 64	Sumata, Hiroshi	56
Robbins, L.L.	77	Suykens, Kim	45
Roberts, Donna	10, 20	Suzuki, Koji	89
Roberts, Jason L.	20	Suzumura, Masahiro	56
Rodolfo-Metalpa, Riccardo	29	Swingedouw, Didier	54, 73
Roevros, Nathalie	45	Takahashi, Taro	22
	33		33
Rogers, Ku'ulei S.		Tan, Adrian	
Rollion-Bard, Claire	76	Tanner, Christina	40
Ross, Pauline M.	18	Tarling, Geraint	40
Russell, Joellen L.	78	Teyssié, Jean-Louis	42, 46, 65, 67
Sabbe, Koen	45	Thatje, Sven	85
Sabine, Christopher L.	15, 37, 48, 78	Thomas, E.	81
Saito, Shu	62	Thomas, H.	82
Sakamoto, Takashi T.	56	Thomsen, Jörn	79
Saphörster, Julia	79	Thorndyke, Michael C.	51, 58, 85
Sartoris, Franz J.	55	Thuesen Erik V.	11, 27, 86
Sasai, Yoshikazu	79	Thuroczy, Charles-Edouard	1 23
Sasaki, Hideharu	79	Tilbrook, Bronte	77
Sasano, Daisuke	62	Todd Martz, Todd	48
Sato, Toru	11, 30	Tokieda, Takayuki	62
Sawada, Yoshifumi	80	Touratier, F.	86
	32	Towanda, Trisha	86
Scheifele, Peter M.		Tréguer, Paul	43
Schmidt, Daniela N.	23, 81		86
Schmidt, Sabine	45	Triantafyllou, G.	
Schneider, Bernd	22	Trübenbach, Katja	84
Schneider, Birgit	81	Trull, Thomas W.	12, 20, 21, 87
Schneider, Kenneth	24	Tsiaras, K.	86
Schulz, Kai G.	12, 31, 32, 41	Tsumune, Daisuke	89
Schulz-Bull, Detlef	73	Tsurushima, Nobuo	56
Schuster, Ute	59	Turley, Carol 13	, 34, 35, 52, 59
Sciandra, A.	25	Turner, Suzanne M,	34
Segschneider, Joachim	73, 82	Tyler, Paul A.	19
Seibel, Brad A.	27	· · · · · · · · · · · · · · · · · · ·	, 44, 51, 57, 59
Seoka, Manabu	80	Ufkes, Judith	26
	00	,	-

Ura, Kazuhiro	80
Van Oostende, Noclas	45
Vance, Thomas	87
Völker, Christoph	57
Voss, Maren	60
Wahl, Martin	41
Watanabe, Yuji	61, 88
Watson, Andew	59
Watson, Sue-Ann	10, 19
Webster, Richard	84
Weinbauer, Markus G.	67
Wesslander, Karin	72
Widdicombe, Stephen	11, 27, 53, 88
Wiebe, Peter	31
Williams, John	57
Williamson, Jane E.	58
Wolf-Gladrow, Dieter	57
Wood, Hannah L.	54, 88
Yamada, Namiha	56
Yamanaka, Yasuhiro	56, 61, 68, 79
Yates, K.	77
Yeats, Philip	39
Yin, Rui	64
Yoshimura, Takeshi	89
Zachos, James C.	16
Zeebe, Richard E.	9, 16, 60
Zöllner, Eckart	23