

## North Atlantic Experiment 1998

RRS Discovery 6 June - 9 July 1998

Coordinator: Dr. Wendy Broadgate Principal Scientist: Professor Peter S. Liss

School of Environmental Sciences
University of East Anglia
Norwich NR4 7TJ
United Kingdom



chlorophyll-a, inset Water-leaving irradiance at 555 nm  $(nL_{\nu}^{555})$  showing coccolith scattering Participants can obtain satellite images from RSDAS via the web site: http://www.npm.ac.uk/rsdas/projects/acsoe/ SeaWiFS 16 June 1998: ACSOE eddy; main picture

## **DISCOVERY 234 CRUISE**

## TABLE OF CONTENTS

		Page no
1.	Acknowledgements	2
	List of Cruise Participants	
3.	Introduction to ACSOE	5
4.	North Atlantic Experiment	6
5.	Modes of Operation	8
6.	Timetable of Events	9
7.	Figure 1 Cruise Track	10
8.	Cruise Reports from Individual Participants	11
	Physical Oceanography - Jane Read	11
	Figure 2 - GPS buoy track	14
	Use of Deliberate Tracers - Phil Nightingale, Fiona Carse and Malcolm Liddicoat	15
	Nutrients, Atmospheric Aerosols, Rain and Iron - Tim Jickells	16
	Figure 4 - Nutrient maps during surveys 1 and 2	20
	Phytoplankton, pigments and volatile Selenium species - Marie Lamren-Larssen	
	and Peter Liss	23
	Figure 6 - Fluorescence and cruise track during Survey 1 and Survey 2	
	Volatile Halocarbons in Air and Seawater - Jon Baker	
	Non-Methane Hydrocarbons in Seawater and Air- Wendy Broadgate	25
	Carbon Dioxide - pCO <sub>2</sub> and TCO <sub>2</sub> - Jonathan Delaney	
	Nitrous Oxide and Methane in the North Atlantic - Thomas Frost	28
	DMS in seawater and air - Adrian Thompson and Laura Cardenas	
	Gaseous peroxides - Andrea Jackson	
	Nitrous oxide, ozone, CN and aerosol surface area - Andrew Allen and Roy Harrison	
	Microzooplankton Grazing and DMS Production - Stephen Archer and Claire Stelfox	
	Size-Fractionated Primary Production and Chlorophyll - Linda C Gilpin	
	Primary Production and Size-Fractionated Chlorophyll - Linda C Gilpin	
	Size-Fractionated Nitrogen Uptake and Ammonium Regeneration –	
	Gwenaëlle Moncoiffé	39
	Bacterial Production and Diversity - Carlos Pedrós-Alió	
	DMS, DMSP, DMSO Speciation and Cycling - Rafel Simó	
	Measurements of DMSP Lyase Activity in Plankton Samples - Michael Steinke	44
	Data Processing and Management - Polly Machin	45
	Technicians Cruise Report - Dave Teare and Jeff Benson	48
9.		
10	0.Samples collected during D234	55
-	Table 1 - Underway Water Sampling Log	
	Table 2 - Log of Events - CTD, Hydrowire, Air, Underway, XBTs	
	Table 3 - CTD stations - date, lat, long, depths	
1	1. Meteorological Data - Weekly Met Plots	
	2. Underway Data - Weekly Plots	
	3.CTD Plots	

## ACKNOWLEDGEMENTS

We would like to thank the master of RRS Discovery, Robin Plumley and his officers and crew for always being helpful and cooperative and ensuring the success of the cruise. Andy Louch and his team at the RVS Operations office, who were friendly and efficient, were invaluable in guiding us through the paperwork in advance of the cruise. As always, the RVS Scientific staff, Dave Teare, Pete Mason, Jeff Benson, Jeff Jones, Rob Lloyd and Martin Bridger were extremely helpful and willing to respond to the many and varied calls for help from scientists, as well as in keeping the instruments running throughout the cruise.

The success of this experiment was critically dependent on finding a stable patch of water to deploy the SF<sub>6</sub> and then to follow this patch for a period of weeks. We are indebted to several people who were not on board the ship, but dedicated much of their time, including many weekends, to the cruise. Steve Groom, Peter Miller and Tim Smyth, from the Remote Sensing Data Analysis Service in Plymouth, provided us with daily satellite images of the area and helpful interpretation of the data. Dave Meldrum, from Dunstaffnage Marine Laboratory, Oban, kindly lent us his GPS buoy and sent us regular faxes of its position throughout the cruise.

We would also like to thank Rosie Cullington for her pre- and post-cruise administrative help and for typing and formatting much of the cruise report.

## LIST OF CRUISE PARTICIPANTS

School of Environmental Sciences, University of East Anglia, Norwich

Peter Liss
Fiona Carse
Wendy Broadgate
Tim Jickells
Adrian Thompson
Jonathan Baker
Michael Steinke
Jonathan Delaney
Marie Hamren-Larsson
Laura Cárdenas

#### Plymouth Marine Laboratory, Plymouth

Phil Nightingale Malcolm Liddicoat Claire Stelfox Steve Archer

Portaferry Marine Laboratory, Queens University of Belfast

Gwen Moncoiffé

Department of Biological Sciences, Napier University, Edinburgh

Linda Gilpin

Southampton Oceanography Centre, Southampton

Jane Read

BODC, Bidston Observatory, Merseyside

Polly Machin

## Department of Marine Sciences and Coastal Management

Tom Frost

Institut de Ciencies del Mar, CSIC, Spain

Rafel Simó Carlos Pedrós-Alió

Environment Centre, University of Leeds

Andrea Jackson

RVS Marine, NERC, Southampton Oceanography Centre, Southampton

Pete Mason SEG
Jeff Jones SEG
Rob Lloyd ISG
Martin Bridger ISG
Dave Teare SIG
Jeff Benson SIG

#### INTRODUCTION TO ACSOE

ACSOE is a 4 year UK NERC Thematic Research Programme to investigate the chemistry of the lower atmosphere (0 - 12 km) over the oceans. The studies aim to bring about a clearer understanding of natural processes in the remote marine atmosphere, and how these processes are affected by atmospheric pollution originating from the continents. This information is vital in understanding regional and global-scale changes in atmospheric chemistry and climate.

#### **ACSOE** consits of three consortia:

- 1. **OXICOA** (OXIdising Capacity of the Ocean Atmosphere), a study of oxidant, radical and related gas chemistry within the clean and moderately polluted marine atmosphere
- 2. **MAGE** (Marine Aerosol and Gas Exchange), a study of aspects of air-sea exchange relevant to atmospheric chemistry and aerosol production.
- 3. **ACE** (Aerosol Characterisation Experiment), a study of gas and aerosol processing through hill cap clouds and subtropical marine stratocumulus.

The North Atlantic Experiment is the final in the series of ACSOE-MAGE experiments.

More information on ACSOE can be found on the web site: http://www.uea.ac.uk/~acsoe/welcome.html

#### THE NORTH ATLANTIC EXPERIMENT

Discovery Cruise D234 6 June - 9 July, 1998

#### Introduction

The exchange of gases and particles across the air-sea interface can have important impacts on the biogeochemistry of both the oceans and atmosphere. For example, air-sea transfer of CO<sub>2</sub> is a large flux in both the natural and man-perturbed cycles of this radiatively active gas. Further, the flux of dimethyl sulphide (DMS, produced by marine algae) from sea to air plays an important role in the production of atmospheric acidity and cloud condensation nuclei (CCN). In the case of transfer of particles, the oceans are a sink for terrestrial mineral dust; a major source of iron which appears to be a limiting nutrient for marine primary production in some important oceanic areas.

The air-sea exchange of trace gases (CO<sub>2</sub>, DMS, and many others) is closely coupled with their production or consumption by the biota in the surface oceans. Hence the processes regulating marine biological production are crucial to understanding, and quantifying, gas exchange. Key factors are the fluxes of the major (N, P, Si) and minor (Fe, Mn, Zn) nutrient elements, some of which in turn are being supplied from the atmosphere, notably N and Fe. Thus, there is an intimate coupling between the inward and outward fluxes of material across the sea surface, which make their combined study, as proposed here, both appropriate and rewarding.

It is well known that concentration distributions of biogeochemical variables in the oceans exhibit marked 'patchiness' on a wide range of temporal and spatial scales. Although much of this variability is intrinsic to the driving forces and biogeochemical processes, 'patchiness' also arises from the way that parameters are observed and measured. Traditionally, biogeochemical observations have been made from ships or fixed platforms. These give insight into variations over the large scale, but only limited information on the nature of processes, since sequential measurements will almost certainly not be in the same body of water biologically (or biogeochemically) speaking. Thus, the nature of many biogeochemical signals is blurred by inadequate deconvolution of the chemical and biological processes from the spatial and temporal variability.

To solve this fundamental problem it is necessary to carry out as near as possible a truly lagrangian experiment, in which the observing system moves with the body of water. Tracking water movement with drogued buoys alone is often unsuccessful, particularly under rough conditions, and gives no information on the effects of small-scale lateral and vertical mixing on the biogeochemical signals. Studies of iron enrichment in the equatorial Pacific (IronEx I and II) and recent studies of bloom development during the PRIME study in the N. E. Atlantic have shown the success and value of deployment of sulphur hexafluoride (SF<sub>6</sub>) as a deliberate tracer for small-scale lagrangian experiments in the open ocean. It is, however, not enough to make detailed process studies at the micro-scale level in such lagrangian experiments if we also require a full understanding of biogeochemical systems. In addition it is necessary to have a proper measure of the meso-scale variability within which to interpret the small-scale observations.

#### **Objectives**

In the ACSOE North Atlantic Experiment we plan to address three main objectives, which are closely integrated.

- 1. To examine the biogeochemical controls of trace gas production, cycling of these gases within the water column and their air-sea exchange by;
- measurement of a number of climatically important gases in air and water,
- examination of factors which control trace gas production in the water column, e.g. phytoplankton species, nutrient status, grazing pressure and bacterial activity,
- estimation of the net fluxes of gases across the air-sea interface,
- construction of trace gas budgets for the mixed layer.
- 2. To investigate the kinetics of air-sea exchange processes by;
- using deliberately added tracers in order to better constrain the fluxes of the gases across the air-sea interface during the experimental period,
- to further advance the state of knowledge concerning the kinetics of gas exchange at sea, particularly in open ocean systems.
- 3. To investigate the coupling of atmospheric inputs to water column biogeochemistry by;
- measuring the deposition fluxes of metals and nutrients to surface seawater,
- investigating the biogeochemical cycling of metals and nutrients in surface seawater, particularly their response to atmospheric inputs,
- examining changes in the plankton community in the patch of water over the period of the experiment,
- looking at short time scale changes in nutrient and trace gas concentrations and biology in surface waters in response to rain events.

North Atlantic Experiment Cruise Plan can be found on the website: http://www.uea.ac.uk/~acsoe/wnortha.htm

## Modes of Operation during the ACSOE North Atlantic Experiment, 6 June - 9 July, 1998

Mode	Measurements	Location	Timing
HYDROGRAPHIC S	SURVEYS	~~~	
Seasoar Survey	Seasoar, T, sal, ADCP, fluorescence, chlorophyll, HPLC, nutrients, trace gases and biological parameters#	Seasoar to 300 m and surface water, occasional CTDs	48 - 60 hr at start, 12 hr at middle and 48 - 60 hr at end of each patch
DAILY ROUTINE			
SF <sub>6</sub> Survey	SF <sub>6</sub> , T, sal, ADCP, fluorescence, chlorophyll, HPLC, nutrients, trace gases and biological parameters <sup>#</sup>	surface water, occasional depth profiles	11.5 hours/dy 1430-0200
Atmospheric	Aerosols and rain (major ions, trace metals, DON, N & S isotopes, organics, CN, surface area), trace gases, NO <sub>x</sub> , SO <sub>2</sub> , O <sub>3</sub> , H <sub>2</sub> O <sub>2</sub> .	Head-to-wind	11.5 hours/dy 0300-1430
	Nutrients, HPLC at 1030, chloro phyll, trace gases and biological parameters <sup>#</sup> , SF <sub>6</sub> profiles	Head-to-wind, surface water, CTD at 1400	11.5 hours/dy 0300-1430
Productivity	<sup>14</sup> C primary productivity (size fractionated?), DOC excretion, <sup>15</sup> N uptake, nutrients, trace gases <sup>#</sup> , DMS and its precursors, bacteria, phytoplankton taxonomy, microzooplankton grazing	Centre of patch pre- dawn CTDs to 300 m, incubations	1 hour/dy 0200-0300
EVENTS			
Rain	Rain collection for major ions, trace metals, DON, N & S isotopes, possible <sup>3</sup> He/SF <sub>6</sub> collection for gas exchange study, bacterial heterotrophic activity	Head-to-wind	whenever it rains!
Diel	Lagrangian water measurements for 48 hours: nutrients, chlorophyll, trace gases and biological parameters*, SF <sub>6</sub> , incubations (trace gase production and biological parameters), Atmospheric measurements (when head-to-wind)	centre of patch next to buoy, CTDs at 2 hourly intervals	48 hours - 1 diel study per patch
CONTINUOUS	met, wind stress, waves, instruments on buoys (pCO <sub>2</sub> )	on ship or buoys	continuous

<sup>&</sup>quot;Trace gases and biological parameters includes DMS, halocarbons, hydrocarbons, N<sub>2</sub>O, CH<sub>4</sub>, <sup>3</sup>He, DMS/DMSP cycling, pH, pCO<sub>2</sub>, TCO<sub>2</sub>, bacterial activity, phytoplankton (abundance, biomass, taxonomic change with time, discrete samples), microzooplankton grazing, some samples will be taken for subsequent Fe and DMSe analyses.

D234 Timetable of events and main sampling modes

Day	JD	Date/Time	General comments	A	Approx time, GMT		
				02:30-04:30	04:30-15:00	15:00-02:30	
1	157	06/06/98	Sailed from Fairlie				
2	158	07/06/98	Passage				
3	159	08/06/98	Passage		$C_{01}$	$C_{02-03}$	
4	160	09/06/98	Passage, hove to				
5	161	10/06/98	Seasoar survey 1	S	S, C <sub>04</sub>	S	
6	162	11/06/98	Seasoar survey 1	S	s	S	
7	163	12/06/98	Seasoar survey 1	S	S	S	
8	164	13/06/98	Deployment of SF <sub>6</sub> , Day 0	S	$C_{05-06}$		
9	165	14/06/98	Day 1	C <sub>07-09</sub>	Α	S	
10	166	15/06/98	Day 2	C <sub>10-12</sub>	Α	C <sub>13</sub> , S	
11	167	16/06/98	Day 3	C <sub>14-16</sub>	Α	B,S	
12	168	17/06/98	Day 4	C <sub>17-19</sub>	A	H <sub>1</sub> , A, S	
13	169	18/06/98	Day 5	$C_{20-21}, H_2$	A	S	
14	170	19/06/98	Day 6, STORM, Diurnal I	C <sub>22</sub>	_	_	
15	171	20/06/98	Day 7, STORM	-22	S	S, C <sub>23</sub>	
16	172	21/06/98	Day 8	C <sub>24-25</sub>	A	C <sub>26</sub> ,H <sub>3</sub> ,S,A	
17	173	22/06/98	Redeployment of SF6	A	A,S,C <sub>27</sub> ,H <sub>4</sub>	C <sub>26</sub> ,113,3,A	
18	174	23/06/98	Diurnal II, transect to Iceland	C <sub>28-29</sub>	$C_{30-33}, H_5, S$	S	
19	175	24/06/98	Heimaey Island, Iceland	€28-29	C <sub>30-33</sub> ,11 <sub>5</sub> ,5	rr	
20	176	25/06/98	transect from/back to Iceland		$H_7$	H <sub>6</sub> H <sub>8-10</sub> , A	
21	177	26/06/98	transect to 59N 21W		A	$A, C_{34}, H_{11}$	
22	178	27/06/98	transect to 59N 21W	C <sub>35-36</sub> , H <sub>12</sub>	S	$C_{37}, H_{13}, S$	
23	179	28/06/98	Day 15 Seasoar deployed	C <sub>38-39</sub>	S,C <sub>40</sub>		
24	180	29/06/98	Seasoar Survey 2	S		S, A	
25	181	30/06/98	Seasoar Survey 2	S	S, A S	S, A S, A	
26	182	01/07/98	Seasoar Survey 2	S, A	S	5, A S, A	
27	183	02/07/98	Survey 2, deep CTD transect	S	S	C <sub>41-43</sub>	
28	184	03/07/98	Deep CTD transect	C <sub>44</sub>	C <sub>45-48</sub>	C <sub>49-50</sub>	
29	185	04/07/98	Final CTDs in study area	C <sub>51</sub>	C <sub>52</sub> , A	C <sub>53,</sub> A	
30	186	05/07/98	Passage	€51	U <sub>32</sub> , A H <sub>14</sub> , A		
31	187	06/07/98	Passage		$C_{54}, H_{16}$	H <sub>15</sub> , A H <sub>17</sub>	
32	188	07/07/98	Passage		H <sub>18</sub>		
33	189	08/07/98	Docked Southampton 2030		1 1 1 8	H <sub>19</sub>	
34	190	09/07/98	Unloading				

Key

S - underway sampling in survey mode

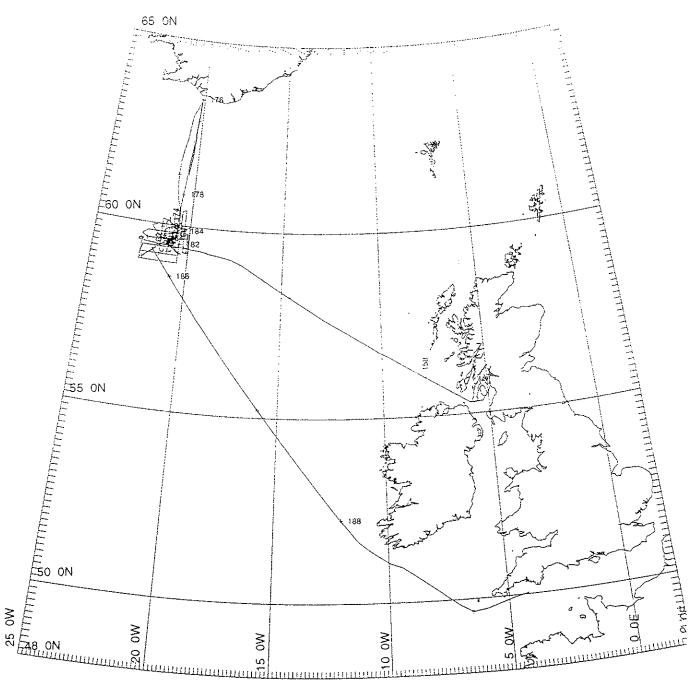
C## - CTD No. ##

A - atmospheric sampling head-to-wind

H - hydrowire sampling B - small boat work

## **CRUISE TRACK**

Figure 1 - Cruise Track of Discovery 234 from Fairlie, Scotland on 6 June to Southampton, England, on 9 July, 1998



AV.

TRANSVERSE MERCATOR PROJECTION

GRID NO. 1

SCALE 1 TO 10000000 (NATURAL SCALE AT C.M.)

C.M. 12W International Spheroid

RRS Discovery 234 Cruise Track

## CRUISE REPORTS FROM INDIVIDUAL PARTICIPANTS

#### Physical Oceanography

# Jane Read Southampton Oceanography Centre, Southampton, UK

The main objectives for the cruise were first, to find and survey an eddy and to locate the centre of the eddy for injection of the tracer gas SF<sub>6</sub>, and second, to investigate the structure and dynamics of the eddy, its temporal variability and the processes involved in its evolution. To achieve these objectives three primary instruments were used to obtain upper ocean currents (ADCP), upper ocean hydrogaphy (SeaSoar) and the baroclinic field (full depth CTDs). Ancillary data were collected as described below.

The cruise began with a SeaSoar and ADCP survey of a feature identified from satellite imagery as an anticyclonic eddy. The data collected confirmed this. Over 3 days a grid pattern was worked with 6 east-west legs 60nm long, 12nm apart and ending with a diagonal line along a Topex/Poseidon satellite track that fortuitously passed almost exactly though the centre of the eddy. From the combination of SeaSoar and ADCP data the centre of the eddy was located to within about a mile, in the northeast quadrant of the survey. ADCP data collected during the ensuing SF<sub>6</sub> injection over a 4 square mile area showed that the tracer was deployed marginally off centre. However, velocity data at 30 and 100m showed there to be sufficient vertical shear in the system to preclude a more exact definition of the eddy centre.

The patch of SF<sub>6</sub> injected into the surface layer remained coherent throughout the next 10 days making any further survey work unnecessary. Further physical oceanography measurements were therefore postponed until the end of the cruise.

During the last week of the cruise a second SeaSoar survey was made. This time the grid was worked in a north-south direction with 8 legs 6nm apart and of varying length, 54-78nm. The close spacing was to enable the data to be analysed for vertical motion where the gradients in all directions need to be calculated. No satellite imagery had been received for over a week and the survey was centred on an estimate of the eddy's position from drifting buoy data. During the SeaSoar survey it gradually became clear that the eddy structure had moved to the east. The extent of the survey was restricted by time constraints and the eddy centre was only crossed on the last leg. It provides an excellent picture of the west side of the eddy but the east side remains unsurveyed.

The SeaSoar survey was followed by a short line of 9 full depth CTD profiles. These were planned to follow a second Topex/Poseidon satellite track that crossed the initial eddy position. During the first part of the cruise the eddy appeared stable and stationary, but during the second SeaSoar survey the eddy centre started translating eastwards and this movement continued throughout the CTD section. By the time the last CTD was worked the eddy centre was some 30nm to the east. The CTD stations therefore did not transect the centre of the eddy as planned but cut across the north west sector.

#### Acoustic Doppler Current Profiler (ADCP)

An RDI 150 kHz hull mounted ADCP was in operation throughout the cruise. During the previous two cruises it had been discovered that one of the four transducers was faulty so had been switched off. This meant that bottom tracking mode was not available, but as the survey region was in deep water and calibration coefficients had been obtained from a zig-zag run on cruise 232, this was not a problem. The ADCP worked well throughout cruise 234. Data were processed using the method established by the IOSDL Marine Physics Group as follows.

ADCP data were logged to a PC using the RDI Data Acquisition System (version 2.48). Two minute ensembles were collected over 64 bins, 8m bin length and 8m pulse length with 4m blank beyond transmit. Since the transducer head is located about 4.5m below the water line the first bin was centred at 13m. Nominal values were provided for speed of sound (1500m/s) and salinity (35.0), with a temperature offset of 45 and temperature scale of 50. Data were corrected for ships heading using the ships gyro. The two minute ensembles were automatically transferred to the ships level C computer and from here they were read into PSTAR data format every 24 hours. The data were first corrected for the ADCP PC clock drift. This ran fast and over the duration of the cruise the correction increased to about 10 minutes. The drift and oscillations of the gyro were corrected by the Ashtech ADU2 ships attitude measurements. Data were calibrated using a misalignment angle of 2.64° and scaling factor of 0.9917. Absolute velocities were obtained by removing the ships speed and direction estimated from the RVS bestnav data stream. Velocities from 101m were inspected daily and showed coherent structure, this was particularly noticeable during the SeaSoar surveys where currents of up to 1 knot were measured and the anticyclonic nature of the eddy was obvious.

#### SeaSoar CTD Measurements

Two SeaSoar surveys were made, the first of 3 days duration at the beginning of the cruise and the second of 3 days 18 hours during the last week of the cruise (table 1). The RVS SeaSoar and associated instrumentation had not been used since the PRIME cruise in June 1996 which might partly explain a number of problems encountered with the data collection.

Initially the SeaSoar failed to work and after 36 hours the fault was traced to a short circuit in the cable termination. Once deployed it was discovered that the level a logging (which averages the 8Hz data to 1 second) was throwing out large quantities of data. This was traced to the 'rate of change of pressure' editing routine, set up for CTD profiling where a much slower rate of change is expected. Replacement level a software was rapidly downloaded. But when data were transferred to PSTAR format for standard processing the salinity data appeared to be rubbish. After some time it was discovered that the level a software was no longer reducing the data to 1 second and 8Hz data were being logged. The time constant needed to compensate for the different response rates of the conductivity and temperature sensors is usually calculated during the data reduction, without this information conductivity was not corrected for time-lagged temperature effects thus generating very noisy salinity data. Averaging of the data showed it to be reasonably good and the old programmes used to process the data before the days of level a logging were resurrected to handle the data. By the time the data were suitably reduced it was found that the salinity data were drifting badly. During the course of the three day deployment the salinity drift amounted to approximately 0.3 psu. Serious fouling and shifts occurred in addition to this. The data are of doubtful quality but may be improved by comparison with the second survey. This sensor (CTD s/n

1055) should not be taken to sea again unless the cause of the problem is established and rectified. The temperature and conductivity sensors were replaced for the next SeaSoar deployment (s/n 2073). The SeaSoar also carried a fluorometer Aquatracker 3 (88-2960-160) and oxygen sensor. The latter was ignored since no oxygen analysis was done during the cruise, hence calibration is not possible and because of the known hysteresis problems of using oxygen sensors on rapidly profiling instruments. Chlorophyll samples were taken during the first SeaSoar survey for post-cruise analysis. From these it should be possible to obtain a calibration for fluorescence to chlorophyll a.

Salinity data collected during the second SeaSoar survey were a marked improvement on the first, once hysteresis had been minimised by increasing the time constant from 0.22 (standard) to 0.5. However offsets of conductivity were frequent, presumably due to biological fouling, and on one occasion were so severe as to necessitate recovery of the SeaSoar so that the conductivity cell could be cleaned. On redeployment salinity showed a significant increase. Such differences will be calibrated out by comparison with underway salinity samples drawn from the non-toxic sea-water supply. Samples were taken at 2 hour intervals on the first SeaSoar survey. These proved inadequate with the large scale drift that occurred and sampling was increased to every hour on the second survey.

#### Full Depth CTD Measurements

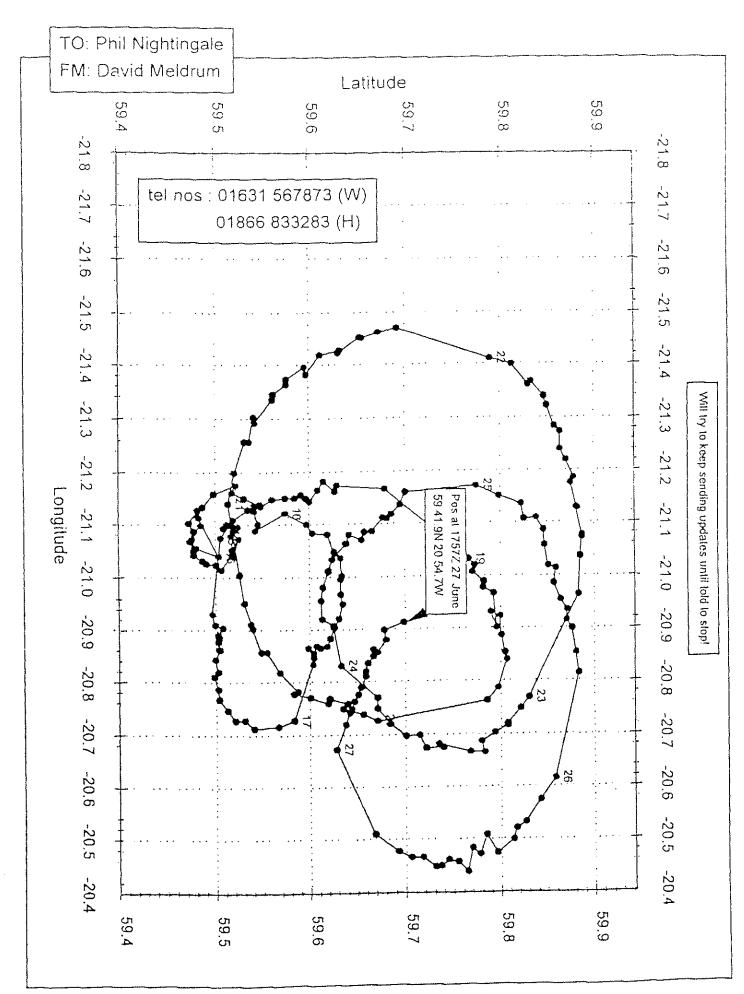
The RVS Neil Brown Mk 3 CTD (s/n 1195) with transmissometer (T1022), nephlometer (SA-244) and fluorometer (Aquatracker 2 SA226) was used to make 9 full depth profiles. Two irradiance meters (nos 1 and 8) used on the shallow casts during the lagrangian study were removed since they were only pressure rated to 1000m. Salinity data were collected from 9 depths; bottom (about 2700-2900m), 2000, 1500, 1000, 500, 200, 100, 50 ,0m.

The 'pinger' (B3) on the first cast gave inadequate returns on approaching the sea bed therefore the cast was stopped about 40m off. The pinger was replaced and the remaining casts were worked to within about 10m of the seabed.

Table 1 SeaSoar deployments

deployment day/time	recovery day time	duration days:hours	distance km
161/0753 (10/6)	164/0600 (13/6)	2:22	644
179/1600 (28/6)	182/0740 (1/7)	2:16	recovered because fouled
182/0900 (1/7)	183/0900 (2/7)	1:0	1252 (combined total)

Figure 2 - GPS Buoy track from Day 0 (13/6/98) of tracer release to Day 14 (27/6/98)



## Use of Deliberate Tracers During the ACSOE D234/98 Cruise

Phil Nightingale\*, Fiona Carse\*\* and Malcolm Liddicoat\*
\*Plymouth Marine Laboratory, Plymouth, UK
\*\* School of Environmental Sciences, University of East Anglia, Norwich, UK

The lagrangian component of this study was provided via a deliberate, controlled release of SF6. A known volume of seawater (2250) litres, saturated with SF6 and 3He, was released at a depth varying between 7 and 9 metres via gravity feed over the rear of the ship for a period of about 10 hours. The release grid chosen was an expanding box with tracks one mile apart and final dimensions of 4 \* 4 miles. In order to avoid loss of tracer to the air while emptying the tank a header tank was used to replace outgoing tracer-tagged water with fresh seawater. The rate of release was increased periodically during the release in order to approximately allow for dilution of the tracer within the tank. Release rates were chosen such that the central one mile of the patch had the greatest concentration of tracer.

Four low profile surface drifting buoys were released during the early stages of the release, one of each corner of the central one mile box. Three of these (RVS owned) were drogued at 20m, the fourth (provided by D. Meldrum CCMS) was drogued at 10m. An in-situ pCO<sub>2</sub> instrument (provided by M. DeGrandpre Univ. Montana) was attached to buoy 1954 at a depth of approx. 16m. This was recovered towards the end of the cruise and will be returned to the US for data downloading.

The tracer patch was successfully tracked for the period of the cruise by using a newly constructed automated system that determined SF6 every 4 minutes. The patch was located after having been hove to during the period of high winds and after an extended period in transit to/from Iceland. Survey periods were used to define the size and movement of the patch and the ship then navigated to the centre of the patch for CTD casts. Samples for the vertical profiling of SF6 were taken from all casts. Samples of seawater were also collected from these casts and stored in cold welded copper tubes for the later analysis of 3He (by P. Schlosser LDEO) in order to derive estimates of gas exchange during the cruise.

The patch was re-infused once using a smaller (1000 litre) tank saturated with SF6 only. The release grid chosen was again an expanding box of final dimensions 2 by 2 miles over a 6 hour period. A check was made that the centre of the patch was indeed being re-infused by monitoring the SF6 in the non-toxic supply from the front of the vessel while the tracer was being released over the stern. The infused patch did indeed cover the centre of the old patch but also extended slightly to the north of the old one. The release was complicated by the movement of the water body at speeds of 1 - 1.5 knots. An attempt was made to allow for this by navigating the ship without attempting to correct for water currents while steaming along the release grid.

A preliminary analysis of the surface data suggests that the tracer in the surface water was sheared from the rest of the water column during the transit to-from Iceland and spread over a fairly large area.

Vertical profiling measurements of Sulphur Hexafluoride

Two sets of 7 profiles were obtained from each of the tracer releases.

#### Tracer I:

The profiles show high concentrations of SF6 (10 - 100 fmol/l) in the top 20m until the 18/6/98. After the storm of 19/6/98, profiles showed mixing to 70m, with surface levels of  $\sim$  10 fmol/l.

#### Tracer II:

The profiles show high concentrations of SF6 (> 100 fmol/l) in the top 20m until the 23/6/98 0600, with lower values (possibly at very edge of patch) until 1330. After the detour to Iceland, the maximum values of SF6 were found at 40m depth, with levels only 2-4 times background in the surface waters.

#### CTD Section:

The first seven stations of the CTD cross-section through the eddy were sampled for SF6. The first 5 profiles showed sub-surface SF6 maxima at 20m, with the final 2 showing minima at this depth. The bottom water samples showed a consistent 0.5 fmol/l SF6 at about 2800m. A contamination test was carried out on all Teflon Niskin bottles in Rockall Trough deep water. This showed negligible SF6 contamination.

Data will be used to calculate vertical diffusivity of the tracer patches in the ACSOE eddy. The CTD section has provided interesting data, particularly the elevated levels of SF6 in the bottom waters.

## Nutrients, Atmospheric Aerosols, Rain and Iron

#### Tim Jickells

School of Environmental Sciences, University of East Anglia, Norwich, UK

#### Water Column Nutrients

Nutrients were analysed on a Skalar San system located in the Chemistry Laboratory. Samples were analysed unfiltered but, in open ocean waters using the standard chemical procedures employed, measured concentrations are assumed to approximate to the dissolved fraction. The instrument was run with three channels analysing for dissolved inorganic phosphorus (henceforth phosphate), dissolved silicon (henceforth silicate) and dissolved nitrate plus nitrite. On one occasion a profile over the upper 200m was analysed for nitrite and concentrations were below detection limit, which is conservatively estimated at  $0.5 \mu Ml^{-1}$ . Hence results for nitrate plus nitrite are subsequently refereed to as nitrate. The same terms are used in the data submitted to BODC.

The instrument set up and analysis procedures were essentially identical to those detailed in previous cruise reports (Sanders R. in Heywood K.B. and King, B.A. 1996 WOCE Section A23 Cruise Report, University of East Anglia Cruise report Series Number 1 and Sanders R. and Jickells T in King B.A. JCR27 Cruise Report in preparation). The instrument generally ran satisfactorily apart from occasional jumps in baseline which occurred simultaneously for

phosphate and silicate, suggesting an occasional electronic fault. This problem made computer data processing inaccurate and the results were therefore derived manually from the chart paper readout. On one occasion a temporary computer fault also required manual processing of data. Comparisons of manual and computer generated results on some runs showed satisfactory agreement, so the different data processing procedures are presumed not to introduce any bias. However, manual processing cannot quantify the extremely low silicate concentrations encountered in surface waters at some stations, and hence the detection limits are different for the two processing procedures. Manually a detection limit of 0.5  $\mu$ Ml<sup>-1</sup> has been used and by computer 0.05 $\mu$ Ml<sup>-1</sup>. The occasions when manual data processing was carried out are identified in the data set submitted to BODC. For nitrate and phosphate no samples that could not be readily quantified were encountered so a detection limit is not reported.

Samples were analysed in batches necessitating refrigerated storage for up to 24 hours, but generally less. The only exception was on the passage leg home when the low sampling density meant samples were stored for up to 50 hours prior to analysis.

#### Accuracy

Standardisation was based on preparation of a stock solution for each analyte in MQ water at the start of the cruise using pre-weighed powders prepared in our home laboratory. Dilute mixed working standards were prepared regularly (usually daily) from these stock solutions.

One set of Sagami nutrient standard reference materials were analysed several times during the cruise. The Sagami silicate standard has a reported concentration of  $100\mu Ml^{-1}$ , well outside the range expected in these waters and outside the range of standards used. This standard was therefore diluted five fold in artificial seawater to give an anticipated concentration of  $20~\mu Ml^{-1}$ . The results in Table 1 are the means of 3 (nitrate and phosphate) or four (silicate) analyses over the period of the cruise and carry the implicit assumption that the standards store satisfactorily once opened.

Table 1 Measured and certified concentrations of Sagami standard reference solutions. All concentrations as μMI<sup>-1</sup>

	Nitrate	Phosphate	Silicate*
Measured	19.4 <u>+</u> 0.1	1.78 <u>+</u> 0.07	20.2 <u>+</u> 0.8
Certified value	20	2.0	20

<sup>\*</sup>After dilution, see text.

The nitrate and silicate results agree well with the certified values. Problems with the Sagami phosphate standard have been discussed previously (Sanders in Heywood 199? and Sanders and Jickells in King 1998) and the results here agree reasonably well with the results we reported previously for other Sagami phosphate standards.

#### Precision

In the early sample analysis runs all samples were run in duplicate. Subsequently sample numbers made this procedure impossible and instead one or more samples were analysed in

triplicate in each analytical run. Based on all these results, the precision of analyses within a single analytical run is estimated at  $\pm 0.1~\mu\text{Ml}^{-1}$  nitrate,  $\pm 0.02~\mu\text{Ml}^{-1}$  phosphate and  $\pm 0.02~\mu\text{Ml}^{-1}$  for silicate. Almost all the replicates were run on low nutrient surface waters. The average of the three deep water precision estimates is essentially identical to those for surface waters. On CTD54, 8 Niskin bottles on the rosette were fired at the same depth, hence providing a check of precision of sampling and analysis. The means and standard deviations for these samples were nitrate  $21.5\pm0.1~\mu\text{Ml}^{-1}$ , phosphate  $1.39\pm0.01~\mu\text{Ml}^{-1}$  and silicate  $35.5\pm0.1~\mu\text{Ml}^{-1}$ .

Long term (between batch) precision was not assessed directly since it seem unlikely that surface seawater samples could be assumed to store without changing concentration. However, an estimate of long term precision can be gained in two ways. Firstly the precision of the re-analyses of the Sagami standard imply precision for high nutrient samples of better than ±5%. Another approach is to consider the analyses of 200m water from the eddy resampled throughout the cruise. As the deepest sample, this seems likely to have been most stable physically and biogeochemically and hence to provide an upper limit on long term precision, though inevitably this is based on deeper water samples with higher nutrient concentrations. Based on 10 re-visitations of the eddy from CTD 6 to CTD 40, long term precision is estimated to be ±6% or better for all three analytes.

#### General Features of the Data

In total about 480 samples were analysed for nutrients from samples collected from underway sampling and CTD casts. The data set has been deposited at BODC. In addition, some extra samples were run at various times as part of incubation experiments and to look for nitrite. Full interpretation of the data set will obviously take some time but some features are already evident.

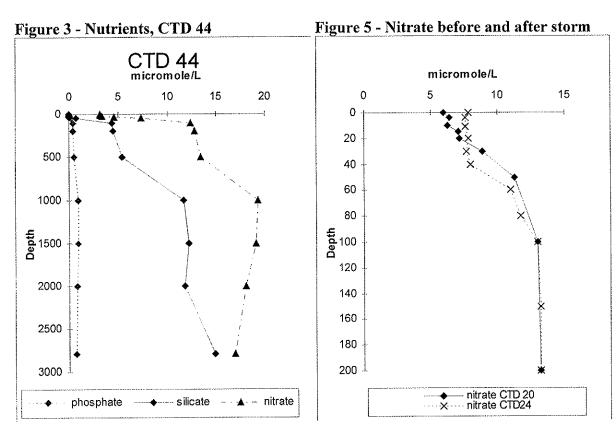
In Figure 3 the nutrient results from the first of the deep CTDs are shown. The profiles show the anticipated low surface water concentrations increasing with depth. Nitrate and phosphate concentrations peak at about 1000m, but silicate peaks in the bottom water suggesting a small component of southern water masses creeping this far north.

The surface water nutrient concentrations were generally low in the area when we first arrived, but far from being totally nutrient depleted. Surface water nitrate concentrations were in the range 5-7.5  $\mu$ Ml<sup>-1</sup>, phosphate 0.3-0.4  $\mu$ Ml<sup>-1</sup> and silicate 0.2-1.2  $\mu$ Ml<sup>-1</sup>. At the time of the first seasoar survey, the eddy structure stands out particularly as a feature with higher surface waters silicate concentrations (Figure 4), presumably due to local upwelling within the eddy.

Throughout the cruise nutrient concentrations appear to decline, though it is difficult to document at this stage from individual profiles since there are marked gradients within the eddy itself. However, comparison of the eddy at the end of the second seasoar survey does imply that nutrient concentrations had indeed fallen, as well as confirming the movement of the eddy. Surface water nitrate concentrations were down to  $3.5-6~\mu Ml^{-1}$ , phosphate  $0.2-0.3~\mu Ml^{-1}$  and silicate concentrations had fallen to effectively zero. The eddy is, however, still identifiable as a region of at least elevated surface water nitrate and phosphate concentrations (Figure 4). At the time of the more detailed second survey there is evidence of another feature with higher nutrient concentrations to the south-west which is also evident in the first survey. There is also a notable region of higher silicate to the north of the eddy which is presumably

associated with upwelling and yet is not associated with higher nitrate levels. An explanation of these features should arise from combining the physical data with both the chemical results and those for phytoplankton growth rates and abundance.

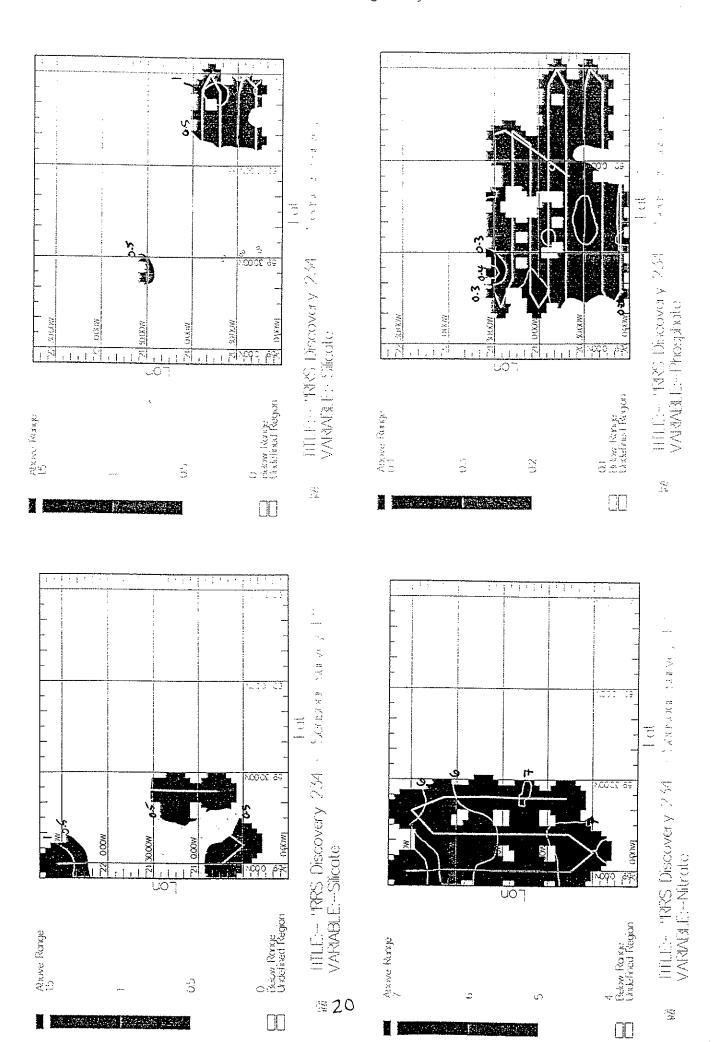
During the cruise, there were several storms passed through the area. These appear to result in important changes in the nutrient profiles due to mixing. In Figure 5 nitrate profiles over the top 200m are shown before and after the first storm that we encountered. The storm has homogenised the water column over the upper 50m and thereby increased nitrate concentrations in surface waters by almost 2  $\mu$ Ml<sup>-1</sup>, with a matching decrease in concentrations in the 30-60m depth range. Silicate concentrations were also increased from 0.3 to 0.8  $\mu$ Ml<sup>-1</sup>. Given that these very low silicate concentrations may be restricting diatom growth, such storm mixing events may be an important process.



#### Iron

Dissolve iron is now recognised as an important nutrient for phytoplankton growth, and hence it is of interest to the ACSOE programme to define its abundance in the waters surveyed. Andy Bowie of Plymouth Marine Laboratory kindly agreed to analyse a small number of samples from the cruise.

Sampling and analysis for iron is a particularly demanding task since the ship is obviously a major potential source of contamination. Surface water samples from within the eddy were therefore collected by hand using the ship's rubber boat about 1 mile from the ship. Sampling from deeper waters was more difficult. The cruise was not originally designed to collect trace metal clean samples and we had not therefore requested clean GoFlo bottles and Kevlar line. We had hoped to use the CTD system with 301 Niskin bottles with teflon coated



springs. Unfortunately upon inspection the coating on the springs had obviously been damaged and there was clear rust staining on the springs making these bottles unsuitable. The RVS technicians were able to provide one spring in much better condition and this was used in a 10l Niskin bottle for which they rigged a short length of plastic coated stainless steel wire mounted on an auxiliary winch to allow sampling of surface waters.

A few deeper water samples were also collected using the CTD bottles with rubber springs. Sampling was arranged so that these bottles had already been extensively used and hence may have been cleaned effectively by use. We will have to await results to see if this procedure has been successful.

Samples were stored frozen after collection for subsequent transport to Plymouth.

#### Atmospheric Sampling

#### (a) Aerosols

Our original plan had been to operate the two aerosol samplers on the bridge deck. Unfortunately communication problems meant that the ship's crew had not been informed of this and several problems ensued, firstly over locating any equipment where it might interfere with the ship's lifeboats and secondly over the absence of a power supply. The former problem was overcome by locating the samplers at the very front of the bridge deck and the latter problem was overcome by the ship's chief engineer agreeing to have power leads led into his cabin and plugged in there. Subsequently earthing problems were found with one of the aerosol samplers and both had to be removed to the laboratory for subsequent repair. After this it was decided to relocate the samplers to the monkey island where more straightforward power supply could be achieved. This transfer was possible only due to the exceptionally calm weather we enjoyed at that time. The samplers were fixed to the aft rail on the port side of the monkey island with power running from sockets at the top of the access ladder. Subsequently this proved a very satisfactory sampling location being high to minimise spray effects and allowing good access even in poor weather. The ship's officers co-operated extremely well in alerting me for rain or changes in apparent or real wind direction.

Subsequent aerosol sampling proceeded well. The two samplers were run with different filtration heads, one with a cascade impactor to provide size segregated samples for subsequent analysis for major ions and trace metals, and the other with a glass fibre filter for collection of material for organic nitrogen analysis and a back up filter to collect SO<sub>2</sub> for isotopic analysis. After several days of sampling, one of the aerosol samplers failed with a suspected burnt out motor. Subsequently the one sampler was run with the different sampling heads on alternate sampling sessions. The winds remained generally from the north throughout the cruise so this alternate sampling strategy did not particularly compromise our primary goals of characterising the aerosol characteristics during the ACSOE experiment. Furthermore the GFC filter will be analysed for major ions so for these (including nitrate, sulphate and ammonium which are of particular interest in the context of this experiment) we will be able to provide as complete a coverage as the sampling regime allowed.

During designated atmospheric sampling periods the ship was turned head to wind, and sampling without the risk of ship contamination was straightforward. On other occasions

during passage or survey work sampling was conducted if it was judged likely that contamination could be avoided, thus maximising the sampling opportunities.

Filters were loaded and unloaded in the clean container and transported to and from the monkey island sealed in plastic bags. Full procedural blanks were run on several occasions. Samples have all been frozen for subsequent return to UEA for analysis. Table 2 provides a complete list of dates when aerosol samples were collected on the cruise.

#### (b) Rain

Holders for rain collectors were mounted on the front rail of the bridge deck. Initially these were mounted upright but subsequently angled into the wind to improve collection efficiency. Two parallel collections were made with one for trace metals and the other for major ions and organic nitrogen. Clean funnels and bottles were prepared for deployment in the clean room and then transported to the bridge sealed in plastic bags. Collectors were in general deployed but covered in anticipation of imminent rain and then uncovered as rain started, assuming the apparent wind direction was such that sampling could be conducted without risk of sample contamination from the ship's funnel. In practice this strategy was not always possible since I was sometimes asleep as rain appeared. In these cases the collectors were deployed as rapidly as possible after rain started and the ships heading allowed clean sampling. The ship's officers were extremely helpful in alerting me to sampling opportunities and to problems arising from changes in wind direction or the ship's heading. During designated atmospheric sampling periods the ship was turned head to wind and sampling without risk of ship contamination was straightforward. On other occasions during passage or survey work sampling was conducted if it was judged likely that contamination could be avoided, thus maximising the sampling opportunities. A complete list of the occasions when rain samples were collected is provided in Table 3, but it should be noted that some of the collected samples may be too small to allow subsequent analysis. In addition several full procedural blanks have been collected.

#### **Table 2 Aerosols Sampling Periods**

```
D1 14/6/98 04.30-15.17
D2 15/6/98 04.25-14.48
D3 16/6/98 04.33-13.40
D4 17/6/98 04.30-20.43
D5 18/6/98 04.30-15.15
D6 21/6/98 04.27-14.47
D7 21/6/98 21.42-22/6/98 08.53
D8 25/6/98 14.30- 26/6/98 03.40
D9 26/6/98 10.30-18.23
D10 28/6/98 16.06-16.40, 21.45 - 29/6/98 08.57
D11 29/6/98 09.31-16.52
D12 29/6/98 19.58-30/6/98 10.35
D13 30/6/98 22.00 - 1/7/98 07.00
D14 1/7/98 14.48-22.38
D15 4/7/98 05.37- 17.51
D16 5/7/98 10.28-19.07
```

#### **Table 3 Rain Sampling Periods**

```
all times GMT unless noted
D1
       6/6/98 21.30(BST)-7/6/98 06.30(BST)
      17/6/98 22.30-18/6/98 06.15
D2
D3
      19/6/98 06.30-10.30
D4
      19/6/98 15.00-21.30
      20/6/98 09.00-13.00
D5
D7&8 23/6/98 07.30-15.00
D10
      24/6/98 21.30-23.15
D11
      25/6/98 00.30-09.00
      4/7/98 06.00-13.00
D12
D13
       4/7/98 16.45-18.42
       5/7/98 12.30-13.30
D14
```

# Collection of Water Samples for Phytoplankton Enumeration, Pigment Analysis and Quantification of Volatile Selenium Species

Marie Hamren-Larssen and Peter Liss School of Environmental Sciences, University of East Anglia, Norwich, UK

#### Phytoplankton Enumeration

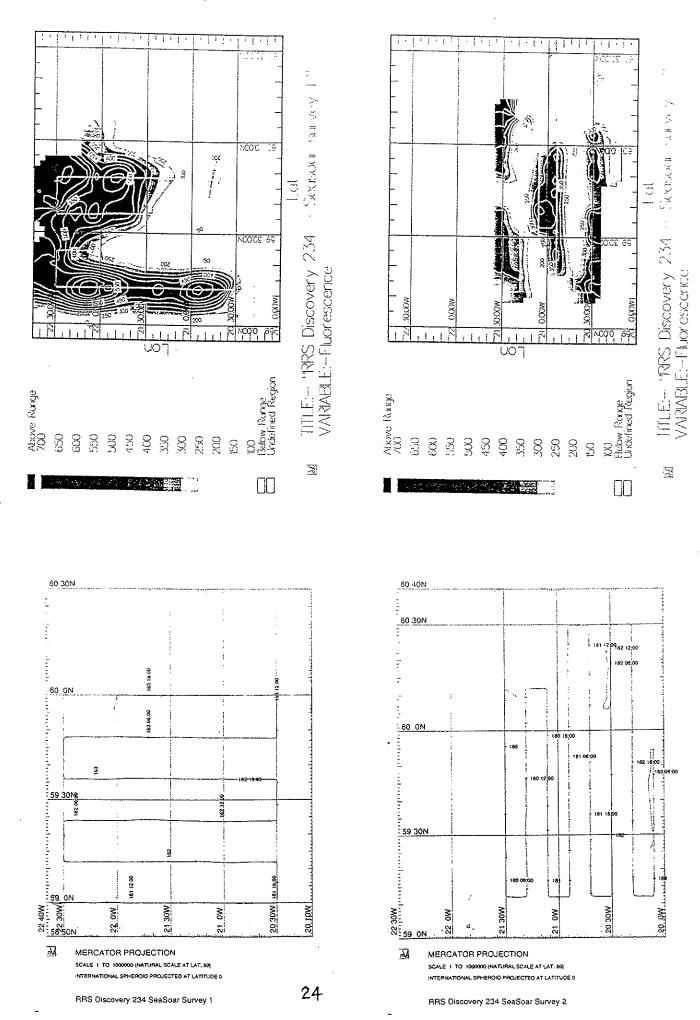
A total of 213 water samples were preserved for subsequent enumeration of phytoplankton species. For each sample two 100ml subsamples were taken, one was preserved by the addition of 2ml of formalin, the other had 1ml of Lugol's iodine solution added. Sample bottles were stored in the dark until counted.

#### **Pigment Analysis**

635 water samples were taken for standard chlorophyll and phaeopigment analysis. Phytoplankton cells were harvested by vacuum filtration of 100ml of sample through a Whatman 25mm GF/F filter. Immediately after filtration the filters were wrapped in aluminium foil and rapidly flash frozen by immersion in liquid nitrogen. Subsequently samples were stored at -80 degrees centigrade prior to extraction and pigment analysis back at UEA. The underway samples were collected from the deck lab (old) non toxic seawater supply direct from the hull and NOT through the debubbling tank. Dave Teare (RVS) informed us towards the end of the cruise that there is a new supply direct from the hull to the TSG and fluorometer which was installed during the refit this winter. After this date (2/7/98) underway trace gas samples, chlorophyll and phytoplankton samples were collected from the outlet of the TSG and fluorometer. Care must be taken when using discrete chlorophylls to calibrate the underway fluorometer before this date as they were taken from different supplies.

In addition 2.15 litre volumes of 160 water samples were filtered through Whatman GF/F filters for subsequent analysis for a range of pigments by HPLC. After filtration the filters were put in plastic vials and flash frozen by immersion in liquid nitrogen. Subsequently the vials were kept at liquid nitrogen temperature until analysed by Stuart Gibb at Plymouth Marine Laboratory.

Figure 6 - Underway fluorescence and cruise track for Survey 1 and Survey 2



#### Volatile Selenium

It has been known for many years that the global budget for selenium cannot be balanced without a flux of some volatile form(s) of the element from the ocean to the atmosphere, with subsequent deposition onto land. However, because of the very low levels of volatile selenium in seawater, it is extremely difficult to prove the emission of selenium from seawater by analytical measurement. Indeed, essentially no measurements of volatile selenium in seawater exist for open ocean situations. In order to try to rectify this deficiency 30 samples of mainly surface seawater were collected using a specially cleaned Niskin bottle attached to a Kevlar line. Within a few hours of sub-sampling of the Niskin into a Schott bottle, the volatile selenium species were extracted from the water using a purge and trap apparatus. The extracted volatiles were trapped using liquid nitrogen and stored in a cryocontainer containing this coolant until analysesd on land. Analysis was carried out by David Amouroux and Emmanuel Tessier using an ultra sensitive GC-MS system in their laboratory at the University of Pau (France).

#### Volatile Halocarbons in Air and Seawater

#### Jon Baker

#### School of Environmental Sciences, University of East Anglia, Norwich, UK

Initial aims were to measure a wide range of volatile halocarbons in air and seawater using a gas chromatograph coupled to a mass spectrometer. Of particular interest was seawater concentrations of CH<sub>3</sub>Br as there is much speculation concerning the source strength of this compound.

Preliminary results suggest that CH<sub>3</sub>Br was undersaturated (30-50%) in surface water of the area sampled. There was very little temporal or spatial variation. Depth profiles showed there to be a surface maximum with concentrations droppping rapidly below the thermocline. Similar trends were observed for CH<sub>3</sub>CH<sub>2</sub>Br.

Other bromine compounds measured (CHBr<sub>3</sub>, CH<sub>2</sub>Br<sub>2</sub>, CHBr<sub>2</sub>Cl) generally had sub-surface maximum concentrations at approximately 40-20 m.

CH<sub>3</sub>I was generally supersaturated in surface waters at approximately 40-400%. Depth profiles demonstrated there to be a maximum at a similar depth to the thermocline on a number of occasions. Similar results were observed for CH<sub>3</sub>CH<sub>2</sub>I, CH<sub>2</sub>CII and CH<sub>2</sub>I<sub>2</sub>.

CH<sub>3</sub>Cl was generally undersaturated in surface waters.

#### Non-Methane Hydrocarbons in Seawater and Air

#### Wendy Broadgate

#### School of Environmental Sciences, University of East Anglia, Norwich, UK

Non-methane hydrocarbons (NMHCs) are important reactive gases in the atmosphere, which provide a sink for the hydroxyl radical (OH) and are key players in the production and destruction of ozone in the troposphere. One reactive hydrocarbon, isoprene (2-methyl 1,3-

butadiene), has strong sources in the terrestrial biosphere, and the ocean as a source has only recently been identified. The mechanism of production of NMHCs in seawater is poorly understood but it is believed to be a combination of both photochemistry and direct emission of NMHCs (and their precursors) from phytoplankton.

Seawater and air samples were collected throughout the cruise and analysed *in situ* by Gas Chromatography ( $Al_2O_3$  PLOT column) and Flame Ionisation Detection. A 1.4 litre sample of seawater was purged with CP nitrogen and the volatile trace gases in the nitrogen stream were passed through several water traps before being cryogenically concentrated and injected into the GC. Approx. 30 aliphatic  $C_2$  -  $C_7$  NMHCs were separated and quantitatively analysed for each sample

Seawater samples were collected from the stainless steel sprung niskin bottles on the CTD, the hydrowire and the ship's underway seawater supply, conincidently with the other trace gases, chlorophyll and phytoplankton samples. The underway samples were collected from the deck lab (old) non toxic seawater supply direct from the hull and NOT through the debubbling tank. This supply was contaminated with some higher hydrocarbons and halocarbons which were not present in CTD samples collected at the same time. Dave Teare (RVS) informed me towards the end of the cruise that there is a new supply direct from the hull to the TSG and fluorometer which was installed during the refit this winter. This supply is of constant flow, is cleaner than the old supply and more reliable. After this date (2/7/98) underway trace gas samples, chlorophyll and phytoplankton samples were collected from the outlet of the TSG and fluorometer.

Air samples were collected via 50 m of 3/4" nylon tube (shared with Jon Baker) mounted on the forward boat deck and pumped into the laboratory via a Charles Austin metal bellows pump at 15 l min<sup>-1</sup>. Upstream of the pump the samples were pumped directly into the analytical system through 1/8" stainless steel tubing. In addition a small number of samples were collected in 3.2 l stainless steel canisters at the inlet of the nylon tube using a battery operated pump.

Underway hydrocarbon water samples were collected routinely during survey mode ie. between ca. 1500 and 0300. Many of the productivity casts were also sampled at ca. 5 depths in the surface 100 m. In addition, samples were collected on the first transects to and from Iceland. Air samples were also collected ca. once daily to confirm the direction of the flux. In all, 101 underway water samples and 81 CTD samples were collected and analysed for NMHCs. CTD casts analysed were numbers 4, 6, 15, 20, 21, 23, 24, 25, 26, 27, 29, 30, 31, 33, 35, 36, 38, 42, 43, 44, 48, 50 and 51. A grazing study (dilution method) was conducted with Claire Stelfox and Steve Archer using samples from 4 m on CTD 51 and 52. The experiment was conducted over 52 hours.

Initial results have only been calculated for three hydrocarbons shown in Figure 7. The isoprene concentration was quite uniform across the patch and the eddy and increased by ca. 30% in the patch over the duration of the experiment. However, on the transect to Iceland we passed through two regions of higher phytoplankton levels which coincided with isoprene increases of a factor of 2. This is consistent with previous data which show a correlation between chlorophyll and isoprene concentration. Ethene and propene increased during the first 7 days of the experiment during sunny, calm weather. Diurnal increases during the day

can be seen for both alkenes during this period. These data support the theory that alkenes are photochemically produced in seawater.

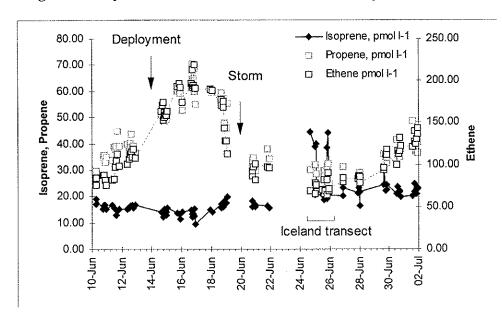


Figure 7 - Surface water NMHC concentrations throughout cruise D234

Depth profiles showed a sub-surface maximum of isoprene. The alkene profiles were of two types (i) exponential decrease from the surface, consistent with photochemical production, and (ii) sub surface maxima in the region of the fluorescence max., perhaps due to the availability of precursors. Two CTD stations (29 and 30) had isoprene levels a factor of 2 higher than all other CTDs in the patch.

## Carbon Dioxide - pCO2 and TCO2

## Jonathan Delaney School of Environmental Sciences, University of East Anglia, Norwich, UK

During the ACSOE research cruise my aim was to measure TCO<sub>2</sub> (total dissolved inorganic carbon) and pCO<sub>2</sub>. The pCO<sub>2</sub> instrument was loaned for this cruise from the University of Bergen and was designed to make routine unattended automated measurements of the CO<sub>2</sub> partial pressure in the ocean surface and overlying marine atmosphere. Unfortunately the pCO<sub>2</sub> instrument ceased to function properly shortly after the start of the cruise, attempts were made to fix it, but as I am unfamiliar with the system I was unsuccessful in obtaining any useful data.

I also had difficulty in using the TCO<sub>2</sub> at the start of the cruise. Generally the problems were electronic faults and programming errors, but one and a half weeks into the research I was in a position to be able to use the TCO<sub>2</sub> instrument. Most of the sampling was under-way analysis from the non-toxic supply onboard, (from a depth of around five meters). This sampling was done both from within the patch and during the detour that was made to Iceland. Where possible I also sampled from the CTD casts that were made.

The results I have obtained are as yet uncorrected and so I will not be in a position to draw any conclusion from the data until my return to UEA. During the research a buoy making pCO<sub>2</sub> measurements was deployed and so I hope to have access to this data in order to compliment my TCO<sub>2</sub> data.

### Nitrous Oxide and Methane in the North Atlantic

### Thomas Frost

# Ocean Research Group, Department of Marine Sciences and Coastal Management, University of Newcastle-Upon-Tyne, UK

The climatically influential trace gases nitrous oxide and methane have been simultaneously measured in both surface and deep water samples of the North Atlantic. Measurement was by a high precision headspace equilibrium gas chromatographic technique, configuration of the system is such that two standard gases and a bow air sample are analysed for every discrete water sample, thus allowing concentrations and saturations to be accurately determined for each sample. A total of 188 underway samples were analysed from the non-toxic water supply and 124 samples were analysed from CTD drops, samples being taken from 23 CTDs. The system performed excellently and only 0.5 days of nitrous oxide data was lost due to an electrical fault with the electron capture detector of the gas chromatograph.

The raw chromatograms have been integrated and the peak areas have undergone preliminary correction for temperature, atmospheric pressure and salinity, although these will be revised when more accurate CTD information is available.

The data will give very detailed coverage of the study area. Upon preliminary investigations into the nitrous oxide data a persistently undersaturated feature was observed during surveys of the study site from the non-toxic supply, as was a subsurface maximum at around 50m.

#### DMS in seawater and air

## Adrian Thompson and Laura Cárdenas, School of Environmental Sciences, University of East Anglia, Norwich, UK

#### Water sampling - introduction

During the daily SF<sub>6</sub> patch surveys surface water was sampled for analysis of dimethyl sulphide (DMS) and its algal precursor, particulate dimethylsulphoniopropionate (DMSP<sub>p</sub>), resulting in a total of 180 DMS samples and around 100 DMSP<sub>p</sub> samples. After each survey, samples were taken from pre-dawn casts for analysis of DMS, DMSP<sub>p</sub>, and dissolved dimethylsulphoniopropionate (DMSP<sub>d</sub>), an intermediate phase in the DMSP<sub>p</sub> to DMS reaction. Along with occasional mid-afternoon casts and casts from aborted diurnals, these data make up a total of twenty depth profiles. Occasionally 10l bottles were used for sampling of DMSe, from which water was also taken for DMS and DMSP<sub>p</sub> analysis. In addition, there was a comparison of different CTD bottles, non-tox/CTD bottle/Niskin bottle, and of two different non-tox supply lines.

Surface water samples were collected from the ship's non-toxic pumped supply (in the Deck Lab) while the ship was in survey mode to determine the spatial and temporal variation in the tracer patch, and to calculate sea-to-air fluxes from the area.

Water samples were collected from bottles on a 'CTD rosette' from a range of depths, usually comprising 5 samples from above the thermocline, and one from below it. Water was collected through a teflon tube into a glass bottle which was rinsed and allowed to overflow to let any degassed water to be replaced with non-degassed water.

The water was analysed immediately (up to 2.5 hours stored in cold dark conditions for some CTD samples) for DMS, by injection through a depth filter into a solid-sorber purge and trap system, followed by flame photometric gas chromatographic detection. The filter was stored in strong alkali (to decompose DMSP<sub>p</sub> to DMS) for analysis at a later date. For waters from depth profile, the purged water sample was quantitatively transferred to a second purge and trap system containing strong alkali (to decompose DMSP<sub>d</sub> to DMS) and analysed using the same gas chromatographic equipment.

Comparison of results, from samples taken from the same CTD bottle, between Rafel Simo, Michael Steinke, and myself showed very good agreement.

(NB. DMS units are given in ngS  $l^{-1}$ . 1ngS  $l^{-1} = 32$ nM)

#### 1. Surface water

Preliminary results suggest that at the start of the study there was an area of high DMS (max = 350ngS l<sup>-1</sup>) at the edge of the survey area, which increased in size to cover the whole survey area over the course of 3-4 days. Following a day of high winds (to force 9) the concentration of DMS dropped to a fairly uniform low, then increased again quickly.

A transect from the patch area to Heimaey was compiled from parts of the two trips north. These show two areas of high DMS: one on the flanks of the Reykjanes Ridge; and one around 62.4N which was associated with increased phytoplankton numbers and hydrocarbon levels.

#### 2. Depth Profiles

Preliminary results suggest that the depth profiles tend to fall into two types: profiles with a subsurface maximum (DMS and DMSP<sub>d</sub>) at 20m or less; profiles with a shallow or even surfacial maximum, and a second maximum at around 40m.

The concentration of DMS seen in the top twenty metres ranged from  $50\text{-}580\text{ngS}\ 1^{-1}$ , and below the thermocline the concentrations were typically around  $10\text{-}20\text{ngS}\ 1^{-1}$ . The range of concentrations of DMSP<sub>d</sub> at the same depths were  $46\text{-}1280\text{ngS}\ 1^{-1}$  and  $10\text{-}40\text{ngS}\ 1^{-1}$  respectively. For both DMS and DMSP<sub>d</sub>, the lowest concentrations were seen after the period of high winds.

#### 3. Measurements of DMS in the marine background atmosphere

#### Introduction

The importance of DMS in the atmosphere has been widely recognised due to its role in climate change. It is a precursor of cloud formation through the production of sulfate aerosol which forms cloud condensation nuclei. These are involved in climate by affecting temperature and radiation (Charlson et al., 1987; Charlson and Wigley, 1994).

The oceanic source of DMS was estimated as about 34% of the total natural sources and about 20% from the total (natural and anthropogenic) (estimate from 1976 in Finlayson-Pitts and Pitts, 1986) making these studies of significance in climate change.

Atmospheric DMS has been measured in the marine background atmosphere on board ships and on shore. Aircraft measurements are also reported in the literature (Andreae et al., 1993; Gregory et al., 1993). Concentrations are below 230 pptv in marine tropical air, (Andreae et al., 1993), below 150 pptv over the Atlantic Ocean (Gregory et al., 1993).

#### 2. Experimental

- 2.1. Site. Measurements of DMS were carried out in the North Atlantic at locations between during the ACSOE (Atmospheric Chemistry Studies in the Oceanic Environment) EAE98 (Eastern Atlantic Experiment 1998) campaign between the 9-6-98 and 6-7-98. Unfortunately, problems with the technique and instrumentation did not allow us to carry out continuous measurements for the duration of the campaign.
- 2.2. Method. Analysis of DMS in the gas phase requires preconcentration of the sample. Gas chromatography is the analytical technique used for further analysis being flame photometry the most commonly used. Interference by oxidants has been observed before, especially from ozone and OH radicals.

In this work the following steps were carried out for the determination of DMS:

- Oxidant removal, by means of a KI trap. This trap was located close to the sample inlet on the mast at about 20 m height. The trap was contained in a cool box inside an ice bath and in series there was a water trap inside an acetone/liquid nitrogen/ice bath (temperature about 20xC). This produced condensation of any water vapour coming from the KI trap (Gregory et al., 1993).
- Preconcentration of the sample. The sample inlet was located at 20 m above sea level and passed through the oxidant and water traps as explained above. Teflon tubing 1/4" O.D. was used for all the sample line. Inside the lab the sample passed through a series of water traps including a glass wool trap (for water drops), an empty glass trap (U-shape) immersed in an ice/water/salt bath and a counter flow system. This one consisted of a nafion membrane inserted in a Teflon tube with a N2 flow of about 400 mL/min going in the opposite direction to the air sample. The air flow was 200 mL/min and it was driven by an air pump located in the outlet of the GC and controlled by a mass flow controller located between the pump and the instrument. Sample was preconcentrated in a loop made of Teflon PFA tubing 1/8" O.D., filled with glass beads and immersed in liquid N2 (temperature about -150xC). Trapping times varied according to the signal obtained (longer times were used when obtaining small chromatographic peaks). Desorbing from this trap was carried out by immersing the trap in boiling water and directing the sample to a second loop made of Teflon PFA tubing 1/16" O.D., filled with Tenax and immersed in an ice/water/salt bath. The presence of a second loop allows to obtain narrower peaks in the chromatograms.
- Sample analysis. Desorption of the sample from the Tenax loop was carried out by inversion in boiling water. By using a 6-port valve the sample could be injected into the GC. For the first period of the campaign a GC VARIAN 3800 equipped with a Pulse Flame Photometric

Detector was used for the analysis. Problems with this instrument made us use a GC SHIMADZU equipped with Flame Photometric Detector available at the site for the DMS water analysis.

- Calibration. Calibration of the detector was carried out by using liquid standards containing DMSP of a known concentration. Reaction of DMSP with NaOH produces DMS which is then trapped by the two loops and analysed by GC. However, this method does not test for any losses occurring in the water traps and/or the sampling line. Ideally, a gas phase calibration is desired in order to test the system fully. For this purpose two gas phase standards were used, a permeation tube belonging to ICM-CSIC with a concentration of DMS of 5.37 ngS ml-1. A second permeation tube was calibrated against the former one and a calibration curve was obtained in order to determine the linearity of the analytical system. This gave a linear regression with a coefficient of determination of 99.5% for levels of sulfur below 16 ng.

Measurements were carried out on days 21-22/6/98, 25-26/6/98 and 2-6/7/98.

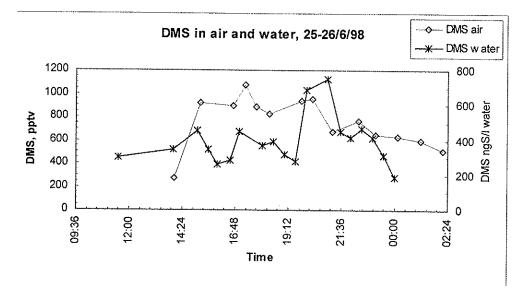
2.3. Results. Atmospheric levels of DMS were between 0.2 and 2 ppbv. Figure 1 shows the results for days 25-26/7 when simultaneous water measurements were performed during the transect South-North to Iceland. Although it is not very clear that there is a direct relationship, a certain similar trend is observed in both sets of data. The levels seem to be higher than levels reported by Andreae et al. (1993) and Gregory et al., 1993. Results presented by Johnson and Bates (1993) at heights below 600 m showed levels below 150 pptv over the North Atlantic and below 104 pptv over the Tropical Atlantic.

Our results seem to be higher than other published data. This could be due to the difference in locations (different productivity areas) among other factors such as meteorology. This must be studied in more detail in conjunction with the measurements carried out in the water.

#### 2.3 References

- Andreae, T.W., M.O. Andreae and H.G. Bingemer (1993) Measurements of dimethylsulfide and H2S over the western North Atlantic and the tropical Atlantic, J. Geophys. Res. 98: 23389-23396.
- Charlson, R.J., J.E. Lovelock, M.O. Andreae and S.G. Warren (1993) Nature, 326: 655-661.
- Charlson, R.J. and T.M.L. Wigley (1994) Sulfate aerosol and climate change, Scientific American, February, p. 48-57.
- Finlayson-Pitts, B.J. and J.N. Pitts Jr., Atmospheric chemistry, John Wiley & Sons, U.S.A., 1986.
- Gregory, G.L., L.S. Warren, D.D. Davis, M.O. Andreae, A.R. Bandy, R.J. Ferek, J.E. Johnson, E.S. Saltzman and D.J. Cooper (1993), An intercomparison of instrumentation for tropospheric measurements of dimethylsulfide: aircraft results for concentrations at the parts-per-trillion level, J. Geophys. Res. 98: 23373-23388.
- Johnson, J.E. and T.S. Bates (1993), Atmospheric measurements of carbonyl sulfide, dimethylsulfide, and carbon disulfide using the electron capture sulfur detector, J. Geophys. Res., 98: 23411-23421.

Figure 8 - DMSwater and DMSair measured simultaneously on the transect to and from Iceland



This was a successful campaign overall during which we achieved all of our original objectives except for a slightly lower survey sampling resolution, and the lack of time to complete high resolution shallow depth profiles.

Many thanks to the crew, officers, RVS, fellow scientists and Peter Liss for making this cruise successful and enjoyable. Particular thanks also go to Wendy Broadgate for an impressive display of scientific, logistical, and administrative juggling.

#### Gaseous peroxides

# Andrea Jackson Environment Centre, University of Leeds, Leed, UK

#### Aim

Gas phase hydrogen peroxide  $(H_2O_2)$  and organic hydroperoxide (ROOH) measurements were made as part of the ACSOE cruise in order to:

- provide an indication of the extent of tropospheric free radical chemistry occurring in the marine boundary layer
- to assist in the interpretation of other chemical measurements
- to provide input to the chemical modelling study

#### Method

Samples were collected during head to wind periods from 10<sup>th</sup> June to 2<sup>nd</sup> July 1998. Measurements could not be made after this time due to a technical failure with the analytic equipment.

Samples were collected by a previously used nebulisation-reflux concentrator for a period of thirty minutes at hourly intervals. This technique provides peroxide extraction from a large

volume of air sample into a small volume of refluxed trapping solution, providing significant preconcentration of the sample, and hence giving good sensitivity.

Samples were analysed immediately by direct injection into a high pressure liquid chromatography (HPLC) system. Analysis is based on the horseradish peroxidase enzyme-catalysed dimerization of p-hydroxyphenylacetic acid. This forms a dimer whose quantitation is based on fluorescence detection. This method is capable of detecting individual ROOH species to a concentration as low as 20 pptv.

#### **Preliminary Results**

Hydrogen peroxide, methyl hydroperoxide (CH<sub>3</sub>OOH) and hydroxymethyl hydroperoxide (CH<sub>2</sub>(OH)OOH) were detected in samples. Hydrogen peroxide was the predominant species being present in almost all samples. It appears that concentrations were mainly in the region 0.25 to 0.75 ppbv but calibrations and corrections are yet to be made. Methyl hydroperoxide was also detected throughout most of the cruise but in much lower concentrations, approximately 0.05 to 0.10 ppbv. This compares well with the few other peroxide measurements made in the marine boundary layer.

No significant diurnal variation in concentration was observed, however, on a couple of occasions a peak in both H<sub>2</sub>O<sub>2</sub> and CH<sub>3</sub>OOH concentration was measured between 8 and 9am.

#### Conclusion

Gas phase H<sub>2</sub>O<sub>2</sub> and ROOH measurements were successfully made during the ACSOE cruise. Data will now be processed and interpreted with respect to other atmospheric measurements made throughout the study. Peroxide measurements will be available in their corrected format as soon as ozone and temperature data can be obtained.

# Continuous Measurements of Nitrogen Oxides, ozone, condensation particle number and aerosol surface area

Andrew Allen and Roy Harrison, Institute of Public and Environmental Health, University of Birmingham, UK

#### Data collection periods

Measurements were made between 15:24GMT June 5th and 18:15GMT July 7th, 1998. No data are available for the following periods:

All measurements: 18:15-18:35 June 17th, 13:35-14:45 June 25th, 10:25-16:35 June 30th. O<sub>3</sub> only: 17:00-23:59 June 9th, 00:00-23:59 June 10th, 00:00-10:15 June 11th, 23:35-23:59 June 14th, 00:00-03:55 June 15th, 07:35-08:55 June 19th, 20:05-23:59 June 21st, 00:00-10:55 June 22nd, 18:45-23:59 June 23rd, 00:00-03:25 16:05-23:59 June 24th, 00:00-11:55 June 25th, 13:45-18:15 July 3rd.

#### Instrumentation

Inlets to all instruments were located on the upper foredeck of R.V. Discovery. The condensation particle counters and Epiphaniometer used inlets of 1.5m lengths of 0.25" diameter copper tubing, while  $O_3$  and  $NO_x$  were transferred to the instruments via a 20m length of 0.25" PTFE tubing. The combined air flow rate through this tubing was 1.7 L min<sup>-1</sup>. All inlets were fitted with small plastic rain and spray covers.

Condensation particle number concentrations were measured using TSI 3022A and TSI 3025A instruments (TSI Inc., St. Paul, Minnesota, USA), which can count particles having aerodynamic diameters of >7 nm (50 % cut-off) and >3 nm (50 % cut-off), respectively. Sampled submicron particles are drawn, at an air flow rate of 300 cm³ min⁻¹, through a chamber containing supersaturated butanol vapour, where they are much enlarged by condensation growth, and into an optical detector where they are counted using either "real-time" or "live-time" modes. The detection ranges of the instruments are from <0.01 particles cm⁻³ up to 10⁶ particles cm⁻³ (model 3022A) and up to 10⁶ cm⁻³ (model 3025A). Mean particle concentrations were calculated from the analog output voltage and recorded at one minute intervals using the Squirrel datalogger (Grant Instruments Cambridge, UK) also used to record the outputs of all other instruments except the Epiphaniometer.

Nitrogen dioxide (NO<sub>2</sub>) and nitric oxide (NO) were determined using an API Model M200 analyser (Advanced Pollution Instrumentation Inc., San Diego, California, USA), which monitors the chemiluminescence produced during reaction of NO with ozone. NO<sub>2</sub> is measured following prior reduction (to NO) in a heated molybdenum converter. This instrument has a detection limit of around 1ppbv.

Ozone (O3) was measured by UV absorption with a Monitor Labs Model 8810 analyser.

Aerosol surface area was measured using a PSI epiphaniometer (Paul Scherer Institute, Switzerland), in which  $^{211}\text{Pb}$  atoms are attached to the surface of atmospheric particles. Subsequent decay of the radioisotopes, measured following collection of the aerosol on a filter, provides an estimate of the Fuchs (or "available") surface area. Data were recorded as 30 minute means using the internal logging system. The design of the instrument inlet allows sampling of particles having diameter  $<5~\mu m$ .

#### Summary of results

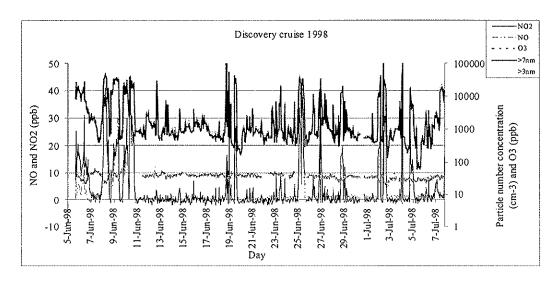
Data are illustrated in Figure 9 (a) (all data for NO,  $NO_2$ ,  $O_3$  and aerosol number concentrations, as 60 minute averages for the entire campaign), and (b) (all surface area data, as 30 minute averages).

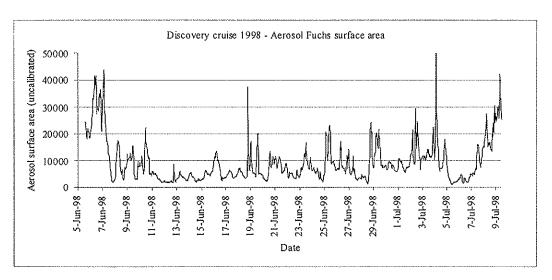
 $NO_x$  concentrations exceeding baseline levels indicate pollution events from sources on Discovery or more distant sources, as do frequent rapid increases in condensation particle number concentrations. Background  $NO_x$  levels were low, and usually close to detection limits although a small increase in background NO concentration occurred after June 22nd. Periods of anti-correlation between  $NO_x$  and  $O_3$  indicate reaction of locally-emitted NO with  $O_3$ . Aerosol Fuchs surface area provides a good indicator of both local aerosol pollution (when increases of surface area occur in line with similar increases of particle number concentration), as well as of the origin of the background air mass (Fuchs surface area is

higher in air masses containing a continental component, when accumulation mode particles having anthropogenic origin provide much of the available surface).

There was no evidence for nanoparticle nucleation during this period. No divergence occurred between the signals of the two particle counters, showing that no production of particles in the range 3-7nm occurred. This is in marked contrast with the coastal environment, where we have previously identified massive photochemical nanoparticle formation, probably driven by oxidation of marine biogenic hydrocarbons emitted by flora within the inter-tidal zone.

Figure 9 (a) Concentrations of NO,  $NO_2$ ,  $O_3$  and condensation particle number (>7nm and >3nm diameter) and (b) Aerosol Fuchs surface area





# Microzooplankton Grazing and DMS Production

# Stephen Archer and Claire Stelfox Plymouth Marine Laboratory, Plymouth, UK

Grazing by microzooplankton on phytoplankton is an important process in the biogeochemistry of both sulphur and carbon in marine surface waters. Microzooplankton are often major consumers of phytoplankton production and therefore carbon and are a key biotic control of phytoplankton population biomass and taxonomic composition. As major consumers of phytoplankton, microzooplankton may play an important role in the fate of phytoplankton DMSP and therefore DMS production.

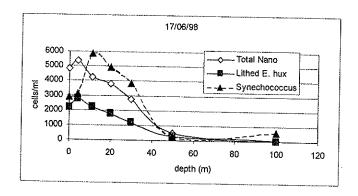
To understand the role that microzooplankton play in controlling phytoplankton populations and in the production of DMS we:

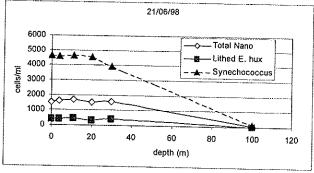
- Collected water samples from the dawn depth profiles to determine temporal and spatial variations in the microzooplankton community structure. Water samples were fixed in acid Lugols, hexamine buffered formaldehyde and glutaraldehyde.
- 2. Carried out grazing experiments using the dilution approach to quantify the temporal changes in microzooplankton grazing pressure on the phytoplankton community within the SF6 patch and outside the eddy.
- Collected water from the dawn depth profiles, CTD transects and during the Seasoar surveys to examine temporal and spatial variations in plankton (picoplankton and nanoplankton) community composition by Analytical Flow Cytometry.
- 4. Carried out incubations in parallel with the dilution experiments from which we hope to budget the fate of DMSP when ingested by microzooplankton and therefore the extent of microzooplankton mediated DMS production.

# Some Preliminary Results:

Phytoplankton populations

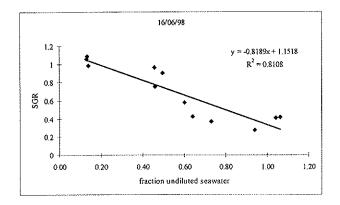
Figure 10. The two plots illustrate the change in depth distribution of the nano- and picophytoplankton before and after the gale force winds

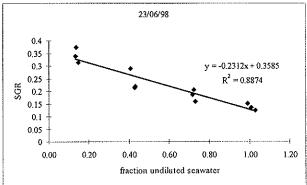




#### Microzooplankton Grazing

**Figure 11**. The two plots are the results of two dilution experiments. They illustrate the lowest and highest grazing pressures measured during the study. Ingestion rates are 13 to 40 ug Chl a  $I^{-1}$  d<sup>-1</sup>, equivalent to a daily turnover rate of 13 % and 56 % of the phytoplankton community in surface waters. The slope of the regression equates to phytoplankton mortality due to grazing (SGR = phytoplankton specific growth rate)





# Size-Fractionated Primary Production and Chlorophyll

# Linda C Gilpin Napier University, Edinburgh, UK

The object of this work was to determine both the rates of primary production and the photosynthetic characteristics of the planktonic assemblages sampled during the ACSOE NAE 1998.

Primary production in the euphotic zone was determined at eight stations using 24hr incubations in an on deck incubator. Water was collected from a predawn CTD cast at 6 depths corresponding to 97, 55, 20, 7, 5 and 1% of surface irradiance - throughout the cruise the depth of the euphotic zone was calculated to be 30 ± 4 m. Six 60 ml samples and two dark bottles were inoculated with <sup>14</sup>C bicarbonate in subdued light and incubated for 24hrs under the irradiance from which the water was sampled. Following incubation, samples were fractionated under minimal vacuum using 5, 2 and 0.2μm polycarbonate membrane filters. Samples of the filtrate were collected for the determination of <sup>14</sup>DOC. Half of the filters were fumed over HCl in order to expel inorganic <sup>14</sup>C such as that incorporated in the liths of coccolithophores while the other filters were not fumed and therefore retained both organic and inorganic fixed carbon. Due to an unfortunate set of circumstances, the particulate incorporation of labelled bicarbonate could not be measured on board. The resulting profiles of carbon uptake per day will be used to determine depth integrated primary production over the euphotic zone at each station.

Attempts to study the diel periodicity in the photosynthetic characteristics of the plankton in a lagrangian mode proved impossible so I undertook two time course experiments with Carlos where a sample was incubated on-deck and subsampled every 4 hours. The photosynthetic characteristics were determined from a photosynthesis versus irradiance (P:I) curve produced following sample inoculation with <sup>14</sup>C sodium bicarbonate and incubation under a range of 24 irradiances using a photosynthetron. Following the incubation, any remaining inorganic <sup>14</sup>C was displaced using acidification and agitation and incorporation of <sup>14</sup>C in the particulate and dissolved components will be measured using a scintillation counter back in the lab.

During the ctd transect, P:I characterisations were made at 5 of the stations. Although these stations occurred at various times of the day, the diel pattern determined from the PI time courses can be used to correct for this. Estimates of column production can be derived from the characteristics by assuming attenuation, chlorophyll distribution and PAR data for the stations sampled.

Size-fractionated chlorophyll measurements were made for each water sample collected during the cruise. Samples were size-fractionated using 5, 2 and 0.2 $\mu$ m membrane filters and the chlorophyll concentration determined fluorometrically following acetone extraction. Results indicate that concentrations  $\leq 1~\mu g \, l^{-1}$  were recorded during the initial stages of the lagrangian experiment. Following our return from Iceland, concentrations up to 1.3  $\mu g \, l^{-1}$  were recorded. Higher values of 2  $\mu g \, l^{-1}$  were observed during the transect and during the aborted diel study.

Duplicate plankton samples from a variety of stations were preserved in 50ml amber bottles using Lugols iodine and Glutaraldehyde solutions.

<u>Cast</u>	<u>Expt</u>	Cast	Expt
5	Si enrichment	39	On deck 7
8	On deck 1	42	PI
11	On deck 2	43	Chlorophyll & taxonomy
15	On deck 3	44	PI time series 2
18	On deck 4	45	Chlorophyll & taxonomy
25	On deck 5	46	PI
29	PI	47	Chlorophyll & taxonomy
30	PI	48	PI
32	PI	49	Chlorophyll & taxonomy
35	PI time series	50	PI
36	On deck 6	52	On deck 8

# Primary Production and Size-Fractionated Chlorophyll

# Linda C Gilpin The Queen's University of Belfast, Portaferry, UK

The object of this work was to determine the species composition, size distribution and photosynthetic characteristics of the phytoplankton assemblages located throughout the LOIS SES working area during the period of the cruise. Stations were sampled across the survey

area to provide spatial coverage of the on shelf, shelf break and oceanic regions; special emphasis was placed on a study of the S and N lines (Table 1).

The photosynthetic characteristics of the planktonic assemblages were determined from a photosynthesis versus irradiance (P:I) curve produced following sample inoculation with <sup>14</sup>C sodium bicarbonate and incubation under a range of irradiances using a photosynthetron. Following the incubation, any remaining <sup>14</sup>C was displaced using acidification and agitation. The incorporation of <sup>14</sup>C in the particulate component will be measured using a scintillation counter back in the lab. A time series P:I study was carried out; a 4 litre sample was incubated on deck with seven subsamples collected between dawn and dusk for the determination of P:I characteristics, chlorophyll concentration and plankton taxonomy.

Primary production in the euphotic zone was determined at four stations using 24hr incubations in an on deck incubator. At station S140 where the euphotic zone was well mixed a sample was collected at 15m from a predawn CTD cast. The depth of the euphotic zone at stations S1500 and N1500 was unknown at the time of sampling so 5m samples were collected. However, at station S700 water samples were collected from a pre-dawn CTD cast at 10 depths selected to represent 97, 55, 33,20,14,7,5,3,2 and 1% surface incident irradiance; the light levels simulated in the incubator. Triplicate 60 ml samples and one dark bottle were inoculated with <sup>14</sup>C bicarbonate in subdued light and incubated at each of the ten light levels in the on deck incubator. Following incubation, samples were fractionated under minimal vacuum using 18, 2 and 0.2µm polycarbonate membrane filters and the particulate incorporation of labelled bicarbonate will be measured using a scintillation counter. The resulting profiles of carbon uptake per day will be used to determine depth integrated primary production over the euphotic zone at each station.

Size-fractionated chlorophyll measurements were made for each water sample collected during the cruise. Samples were size-fractionated using 18, 2 and 0.2µm membrane filters and the chlorophyll concentration determined fluorometrically following acetone extraction.

Duplicate plankton samples from each station were preserved in 50ml amber bottles using Lugols iodine and Glutaraldehyde solutions.

# Size-Fractionated Nitrogen Uptake and Ammonium Regeneration

# Gwenaëlle Moncoiffé Queen's University of Belfast, Portaferry, UK

The main objective of this work was to study daily and diel changes in some aspects of nitrogen cycling by microplankton community in the euphotic zone during a lagrangian experiment. Nitrate uptake and ammonium uptake and remineralisation were measured using stable isotope nitrogen-15 techniques. The rates of uptake were assessed in two size-fractions:  $<200 \mu m$  and  $<5 \mu m$ . Samples were collected under three modes: daily pre-dawn CTD casts, diel CTD casts and CTD casts along a cross-section of the eddy studied during the lagrangian experiment.

Under the daily pre-dawn CTD mode, samples were collected from 6 depths within the euphotic zone (estimated from the depth of the 1% isolume). Profiles of light penetration did not change significantly over the course of the experiment and the depths sampled were: 1-2 m (97% of surface irradiance), 4 m (55%), 11 m (20%), 17 m (7%), 20 m (4.5%) and 30 m (1%). Aliquots of water from each depth was prefiltered through 200 μm mesh and separated into two carboys. Each carboys were spiked with small amounts of sodium <sup>15</sup>N-nitrate and <sup>15</sup>N-ammonium chloride respectively. Spike concentration were calculated so that nutrient enrichment from <sup>15</sup>N addition would not exceed 10% of ambient concentration. For nitrate addition, this amount was estimated from the *in situ* profiles obtained on board by Tim Jickells. Measurement of *in situ* ammonium concentration was not available during the cruise, however literature data from previous experiments carried out at this time of year in this area have shown that ammonium concentration is generally in the nanomolar range in the euphotic zone. As a result minimum additions of 0.02 to 0.04 μM were selected.

Immediately after spike addition, aliquots of water were subsampled and analysed for determination of initial nutrient concentration, nutrient isotopic enrichment and particulate organic nitrogen (PON) concentration. The remaining of the spiked water was incubated on deck under simulated *in situ* light condition. Incubation period was 6 hours for ammonium-spiked samples and 12 hours for nitrate-spiked samples.

Seven experiments were carried out under the pre-dawn CTD mode on 14/6, 15/6, 16/6, 17/6, 21/6, 27/6 and 28/6/98. One of this experiment (27/6) was carried out on the way down from Iceland and was outside the SF6 patch.

From these experiments, profiles of size-fractionated ( $<200 \mu m$  and  $<5 \mu m$ ) particulate organic nitrogen, of size-fractionated nitrate and ammonium uptake, and of ammonium remineralisation rates by the microplankton community will be obtained. In addition, *in situ* profiles of ammonium concentration in the upper 30 m of water column will be obtained using an indirect sensitive method based on the isotopic composition of the ammonium pool immediately after <sup>15</sup>N-spike addition.

Under the diel CTD mode, sampling frequency from CTD cast at the centre of the SF6 patch was scheduled for every 6 hours in order to carry out four experiments of four hours incubation period daily. Samples were collected from two depths (surface and 20 m) and processed as above. Due to adverse conditions, all attempts to carry out a complete diel study during the cruise failed and <sup>15</sup>N data from such experiments will only be available from CTD casts 29 (04:00) and 31 (08:00) on 23 June.

Five <sup>15</sup>N experiments were carried out along the CTD line across the eddy area between 02 and 04 July. These were made on alternate CTD casts (42, 44, 46, 48 and 52) with water sampled from the surface and 20 m. Because samples were drawn from different water masses and at different time of the day, incubation period was extended to 24 hours so as to allow meaningful comparison between stations. Extra volume of water was also collected at the surface from the first CTD station in order to carry out a 24 h time-course experiment and correct for potential losses of <sup>15</sup>N tracers associated with long incubation periods.

Finally, a silicate enrichment experiment was conducted on 13 June 1998 at the end of the first Seasoar survey prior to SF6 release with water collected at a station located along the southern edge of the eddy chosen for SF6 release. The decision to carry out such experiment

was made after surface mapping of nutrient concentration during the Seasoar survey revealed that silicate concentration was below 1 µM across the area surveyed and hence could have been an important factor limiting diatom growth and possibly nitrate uptake in the area. Water for the experiment was drawn from 10 m (CTD 5) and distributed into four 4.5 L incubation bottles. Two bottles received 3 µM of silicate and <sup>15</sup>N-nitrate tracer (10% of ambient concentration) and two were used as controls spiked with <sup>15</sup>N-nitrate but not silicate addition. The experiment was stopped after 24 h. In addition to nitrate uptake rates, subsamples for nutrients analysis (Tim Jickells), <sup>14</sup>C uptake (Linda Gilpin), Chl a (Linda Gilpin and Claire Stelfox) and analytical flow cytometry/phytoplankton composition (Claire Stelfox and Steve Archer) were drawn either on a time-course basis or at the beginning and end of the incubation.

The time-course of nutrient concentration in the control and silicate-enriched bottle suggested that silicate addition had no apparent effect on the phytoplankton community over 24 h; while the evolution in nitrate and phosphate concentrations indicated that considerable uptake was taking place over the 24 h incubation period changes were not significantly different in the silicate-enriched sampled compared to the control. Furthermore, the time-course of silicate concentration indicated that the addition of 3  $\mu$ M of silicate did not trigger any uptake response by the phytoplankton community suggesting that either diatoms were absent from the mixed layer or that their response-time to silicate enrichment was longer than 24 h. The outcome of this experiment may prove useful in interpreting some of the development in the nutrient field distribution reported by Tim Jickells (and perhaps phytoplankton dominance?) after the storm during the second half of the cruise; following an initial increase of surface silicate concentration due to a deepening of the mixed layer, silicate decreased down to level below analytical detection over most of the survey area indicating that diatom growth had taken place after the storm.

Time course of nutrient concentration (nitrate, phosphate and silicate) in control and in water sample enriched with 3 µM of silicate during 24 h incubation period.

# **Bacterial Production and Diversity**

Carlos Pedrós-Alió

Departament de Biologia Marina i Oceanografia, Institut de Ciències del Mar, CSIC, Barcelona, Spain

#### **Objectives**

- 1. To determine genetic diversity of planktonic microorganisms (Bacteria, Archaea and Eukarya) along a Lagrangian time series and through several water masses (eddy transect).
- 2. To determine abundance and heterotrophic production of bacterioplankton.

#### Methods

1. Collect microbial biomass on 2 μm polycarbonate filters and on 0.2 μm Sterivex filters. Extract DNA, PCR for 16S rRNA genes with Bacterial, Archaeal and Eukaryal

- primers, and carry out denaturing gradient gel electrophoresis (DGGE). On selected samples clone PCR products and sequence.
- 2. Fix samples in paraformaldehyde plus glutaraldehyde and freeze in liquid nitrogen. Determine pico and nanoplankton abundance by flow cytometry. Fix samples in formalin, filter through 0.2 μm polycarbonate filters and stain with DAPI. Determine prokaryotic abundance and cell volume by image analysis.
- 3. Incubate samples with tritiated leucine and tritiated thymidine. Stop incubations with TCA, centrifuge, rinse with TCA and add scintillation cocktail. Determine radioactivity incorporation by liquid scintillation counting. Determine conversion factors between leucine and thymidine incorporation and bacterial heterotrophic production in filtration-dilution cultures.

## Summary of work

## A. Samples processed

- 52 Filters (0.2µm) for bacterial numbers and cell volume by epifluorescence
- 325 Samples fixed for picoplankton abundance by flow cytometry
- 218 Leucine incorporation determinations
- 275 Thymidine incorporation determinations
- 72 Filters (0.2µm) for Archaeal and Bacterial genetic diversity
- 72 Filters (2 µm) for Nanoplankton genetic diversity
- 72 Filters (GF/F) for >2μm and <2μm chlorophyll determination
- 12 Filters for Coccolithophorid diversity by SEM

## B. Lagrangian time series

Ten (6 consecutive) daily vertical profiles of picoplankton abundance, bacterial heterotrophic production and pico- and nanoplankton genetic diversity.

One profile outside eddy and one prior to seeding of patch with SF6.

## C. Transect across eddy

Eight vertical profiles of picoplankton abundance, bacterial heterotrophic production (6 depths each) and pico- and nanoplankton genetic diversity (0m and 500m).

#### D. Diurnal cycles

Two experiments with on deck incubations of surface water. Seven points covering 24 hour periods taken in each one for bacterial heterotrophic production. Bacterial abundance at the beginning and end of incubation (in collaboration with Linda Gilpin).

#### E. Miscellaneous

Two experiments to determine conversion factors from leucine and thymidine incorporation to bacterial heterotrophic production.

One experiment to determine linearity of incorporation of thymidine and leucine with time. Three experiments to determine the effect of chloroform, n-serve and DMDS on bacterial heterotrophic production (with Rafel Simó).

# DMS, DMSP, DMSO Speciation and Cycling

Rafel Simó Departament de Biologia Marina y Oceanogrfia Institut de Ciencies del Mar, CSIC, Barcelona, Spain

## **Objectives**

- 1. To determine the speciation of DMS (including DMSPd, DMSPp, DMSOd and DMSOp) along a Lagrangian time series.
- 2. To estimate the biological production and consumption rates of DMS and DMSP along a Lagrangian times series.
- 3. To study the short-term variability of both the speciation and the transformation rates in a 24 h time series.

#### Methods

- 1. Collect water samples from CTD bottles in glass flasks, and run immediate DMS analyses in GF/F filtrates by purge&cryotrap gas chromatography.
- 2. Hydrolyse the sparged filtrates with NaOH for at least 6 h. Determine DMSPd as the evolved DMS. Neutralise the sample and determine DMSOd by borohydride reduction to DMS.
- 3. Hydrolise the GF/F filters with NaOH for aat least 12 h. Determine DMSPp as the evolved DMS. Neutralise the sample and detrmine DMSOp by borohydride reduction to DMS.
- 4. Incubate whole seawater samples in dark 2-L bottles with and without addition of dimethyldisulphide (DMDS) as an inhibitor of DMS consumption. DMS production rate is estimated from the slope of the time course of DMS concentration in the inhibited sample. DMS consumption rate is calulated as the difference of the slopes of the inhibited and the control samples. DMSP consumption rate is estimated from the slope of the time course of DMSPp+DMSPd concentrations in the control sample.

### **Summary of Work**

### A. Lagrangian time series

DMS speciation in 11 surface water samples out of a 15-day time series. Most from pre-dawn casts. Production and consumption rates in 7 of the water samples mentioned above.

## B. Non-lagrangian waters

DMS speciation in 6 surface water samples outside the SF6 patch, or even outside the eddy. Production and consumption rates in 5 of the samples mentioned above.

## C. Diurnal cycle

DMS speciation and production&consumption rates in on-deck incubated water. Seven time points covering a 24 h period. DMSP and DMSO concentrations from this experiment to be determined in Barcelona.

### D. <u>Miscellaneous</u>

Incubation experiment with addition of N-serve -an inhibitor of mono-oxigenase in nitrifying bacteria- to study the potential role of these bacteria in DMS consumption and DMSO production. DMSP and DMSO to be analysed in Barcelona.

Determination of DMSOd in Michael Steinke's experiments of DMS photochemistry. Most of DMSO concentrations to be determined in Barcelona.

#### Main Achievements

In the lagrangian time series, DMS concentrations ranged 2.8-9.8 nM; DMSPd 6.9-24.5 nM; DMSPp 25.3-174.4 nM; DMSOd 2.1-11.5 nM; and DMSOp 1.9-33.4 nM.

DMSP consumption rates ranged 8-78 nM/d; DMS production 1-19 nM/d; and DMS consumption 0-7 nM/d. These rates were quite variable, showing a dynamic coupling/decoupling of production and consumption processes.

This dynamics was also evident in a 24-h basis. Interestingly, biological transformation rates predicted very well the short-term measured variability of DMS in the diurnal cyle experiment.

# Measurements of DMSP Lyase Activity in Plankton Samples

# Michael Steinke School of Environmental Sciences, University of East Anglia, Norwich, UK

#### **Objectives**

Is algal DMSP lyase present during a coccolithophorid bloom, or are bacterial pathways the major source of DMS?

## **Brief Description of Methods**

On board experiments were carried out to investigate the DMSP lyase activity in particulate samples from various depths. Water samples were filtered (about 2 litres per enzyme test) and filters suspended in a buffer before particles were homogenised with an ultrasonic probe. The resulting crude extract was analysed for enzymatic production of dimethyl sulphide (DMS) from exogenous dimethyl sulfoniopropionate (DMSP) in an *in vitro* assay.

A gravity size-fractionation of particles (18 (10)-200  $\mu$ m, 5-18 (10)  $\mu$ m, 2-5  $\mu$ m and 0.2 to 2  $\mu$ m) was used to investigate relative enzyme activity in phytoplankton size classes compared to bacterial DMSP lyases.

## Sampling

Water samples (56 enzyme tests total) were taken from the surface using either bucket sampling (4), the non-toxic seawater supply system (4), Niskin bottle samples (SEB; 2 tests), CTD surface bottles (16) or within the upper 100 m from CTD depth profiles (30). Most of the CTD depth profiles (4 out of 6) were taken during a transect from CTD #42 to CTD #50. An intensive surface sampling was carried out during a passage from and towards Iceland which includes a comparison between coastal and oceanic waters.

Additionally, water samples were taken for the isolation of various plankton organisms, which will later be tested for DMSP lyase activity.

## **Preliminary Results**

DMSP lyase activity was present in particles >2 µm in situ. Most of the previous research on enzymatic DMS production focused on isolated cultures of bacteria, phytoplankton and benthic macroalgae. Only one study investigated the DMSP lyase activity during a *Phaeocystis* bloom in coastal waters of the North Sea (Stefels, J. et al. 1995. Marine Ecology Progress Series 123:235-243). Therefore, this is the first evidence that algal DMSP lyases may be important for the production of DMS within an oceanic coccolithophorid bloom.

Snap-freezing of filters in liquid nitrogen followed by storage for several weeks at -80°C resulted in only minor loss of DMSP lyase activity (less than 10 %).

#### Outlook

Activity measurements of samples brought back from the field is underway including a partial characterisation of DMSP lyases. Nutrient and plant pigment data, bacterial counts, bacterial metabolic activity and phytoplankton speciation will be needed to discuss the results and to highlight plankton groups most probably involved in enzymatic DMS production.

## **Data Processing and Management**

Polly Machin Bidston Observatory, Birkenhead, UK

#### Aims

- To obtain a complete record of all sampling events on the cruise
- To monitor the performance of the underway meteorological data
- To assist with SeaSoar and ADCP processing
- General watchkeeping/ sampling duties as required.

#### 1) Data Logging

Thanks to the efforts of everyone, a complete record of all sampling events was kept. These included:

54 CTD casts
11 (successful) XBT profiles
12 bucket samples
19 ten litre water bottle deployments
381 samples taken from the non-toxic supply x air samples
16 aerosol samples
11 rain samples

These will be used to create a relational database which is effectively a record of everything (scientific!) that went on during the cruise. This can then be used to answer questions such as .

- what other measurements were made at the same time as my non-toxic sample?
- what measurements were made within 5 minutes of my air sample?
- what was the daily integrated solar radiation for the Lagrangian experiment prior to the first storm?
- how much did the ship drift during each of the deep CTDs?
- what historical data is there in this geographic area?

A complete listing – in time order – of all sampling events is given in Appendix 1.

## 2) Meteorological Data

Checks were made on the continuously recorded meteorological data as follows:

PAR and solar radiation data were checked to see that output was sensible.

Air temperature was checked against the bridge sensors and found to correlate well.

Barometric pressure was checked against the bridge barometer and found to read consistently low by 2-3 millibars. This is partly due to the fact that the bridge barometric pressure is corrected to a height of 10 metres above sea level.

Relative wind velocities were converted to absolute and checked against the 6 hourly observations made the ship's officers. Approximately 80% of the wind speeds agreed to within 5 knots and about 70% of the wind directions agreed to within 25 degrees. There were a few instances when the estimated and measured wind directions differed wildly, but in all cases this was at low wind speeds and probably due to the wind swirling. The anemometer was noted to be slow to respond to changes in wind direction. This will be investigated in more detail at BODC.

The assistance of the Captain and Officers of RRS Discovery in making their 6 hours met.obs. books available is gratefully acknowledged.

# 3) Where were those CTDs ??

CTD1,CTD2,CTD3	59 20 North, 19 West	outside the eddy
CTD4	59 North, 22 30 West	outside the eddy
CTD5	59 15 08 North, 20 32 45 West	outside
CTD6	59 30 20 North, 21 07 55 West	inside
CTD7,CTD8,CTD9	59 30 19 North, 21 07 47 West	inside
CTD10,CTD11,CTD12	59 28 44 North, 21 04 50 West	inside
CTD13	59 32 50 North, 21 03 34 West	outside
CTD14,CTD15,CTD16	59 33 59 North, 21 06 54 West	outside upwelling region
CTD17,CTD18,CTD19	59 33 45 North, 20 46 07 West	inside
CTD20,CTD21,CTD22	59 31 26 North, 21 00 01 West	inside
CTD23,CTD24,CTD25	59 33 12 North, 21 01 36 West	inside
CTD26	59 34 47 North, 21 08/38 West	edge
CTD27	59 53 North, 21 15/23 West	edge
CTD28,CTD29,CTD30	59 54 19 North, <b>2</b> 0 44 30 West	inside
CTD31,CTD32	59 52 03 North, 20 34 17 West	edge
CTD33	59 50 52 North, 20 39 35 West	edge/outside
CTD34	61 11 50 North, 20 31 14 West	outside
CTD35,CTD36	60 25 54 North, 20 39 37 West	inside upwelling region
CTD37	59 45 38 North, 21 West	inside
CTD38,CTD39	59 48 26 North, 20 57 53 West	inside
CTD40	59 49 57 North, 20 41 47 West	edge/outside
CTD41,CTD42	59 59 56 North, 20 04 02 West	feature to North
CTD43	59 53 04 North, 20 21 48 West	outside
CTD44	59 46 22 North, 20 39 11 West	edge
CTD45	59 39 41 North, 20 56 58 West	outside
CTD46	59 32 42 North, 21 13 48 West	outside
CTD47	59 25 43 North, 21 31 24 West	salty feature
CTD48	59 18 45 North, 21 48 21 West	outside
CTD49 /	59 11 38 North, 22 05 20 West	outside
CTD59	59 04 31 North, 22 21 53 West	outside
CTD51,CTD52	59 20 22 North, 21 44 43 West	outside
CTD53	59 20 30 North, 21 55 28 West	outside
CTD54	54 24 30 North, 14 50 10 West	outside

N.B. The position of the CTDs relative to the patch/eddy was determined by the concentration of SF<sub>6</sub> in most cases. In CTDs 42 to 54, it was determined with reference to the preceding SeaSoar survey and in CTDs 14,15,16,35, and 36 it was determined by examining the silicate concentrations.

# **Technicians Cruise Report**

# Dave Teare and Jeff Benson RVS Operations, SOC, Southampton, UK

#### **CTD**

- 1) A total of 54 CTD casts were completed during this cruise. The sensor configuration was as follows:
  - Aux 1) Transmissometer
  - Aux 2) Fluorimeter
  - Aux 3) Down-welling irradiance
  - Aux 4) Up-welling irradiance
  - Aux 5) Nephelometer
  - Aux 6) Not Used
- Transmissometer serial number T-1022D was used from cast 001 through cast 009; transmissometer serial number T-1018D was used from cast 010 onwards. The change in sensors was for the purpose of investigating viability of the second transmissometer. Data from sensor T-1018D was consistent with the nephelometer for the duration of the cruise.
- Fluorimeter serial number SA-226 was replaced with fluorimeter serial number SA-234 beginning with cast number 010 because of consistently low voltage readings from sensor SA-226, (less than 0 VDC below the chlorophyll peak.)
- 4) Irradiance sensors removed for deep casts, number 041 through number 050, and number 54.
- A small quantity of water was observed in the oxygen sensor housing after cast 33, and the amount of water observed increased after the deep casts, (041 through 050). The oxygen sensor and 'O' ring were changed before the cruise due to water ingress; the housing itself may be at fault.
- 6) Continuous power module for the rosette failed during cast number 002, and was replaced with the power module for the duration of the cruise.
- 7) All other instruments/sensors functioning properly.

#### **ADCP**

1) With the exception of the three-beam operational status (beginning with cruise DY232), no problems experienced.

#### **SURFMET**

1) The surfmet system consisted of the following sensor suite:

- A) starboard and port 2-pi par light meters
- B) starboard and port total irradiance light sensors
- C) air temperature
- D) relative humidity
- E) barometric air pressure
- F) wind speed and direction
- G) transmissometer/fluorometer
- H) housing/remote surface sea water temperature
- I) conductivity/salinity for surface sea water
- 2) Both the transmissometer and fluorometer were removed and cleaned, with open air values taken from each, once per week at the beginning of the cruise and for the first two weeks into the cruise. Thereafter the sensors were cleaned on a schedule of twice per week to the end of the cruise.
- 3) Port and starboard 2-pi PAR light meters reading different values, with an offset of approximately 1.5 between the two sensors. Calibration data not available onboard.

## **Fixed Equipment**

- 1) Chernikeef log repeaters reported no working problems.
- 2) Shipborne Wave Recorder would periodically report transfer byte errors and stop logging. Logging was successfully resumed after each error.
- Simrad EA-500 echosounder not working on both port and starboard hull transducers. Both 10kHz and 12 kHz frequencies were tested with similar non-functioning results. PES fish deployed mid-cruise and functioned with no problems to the maximum depth recorded of 2800+ metres.
- 4) A total of 11 XBT's were launched on the cruise; all were T-5 probes and were transmitted to GOES. Not included in these launches were a total of 5 failures, due all or in part to rough weather, excessive ship speed, or probe malfunction.

#### **PORTASAL**

- 1) The portasal was installed in the constant temperature lab or the darkroom for the cruise and reported no problems with the exception of occasional air bubbles becoming trapped in the far upper end of the conductivity cell. Light, gentle blowing into the flush port removed the bubbles.
- 2) A total of 112 CTD/TSG salinity samples were taken on this cruise.

#### **SEASOAR**

- 1) The Seasoar system consisted of the following sensor suite:
  - A) Mark IIIB CTD

## B) Fluorimeter

The installed CTD, serial number 01-1055, was found to have a drifting conductivity sensor during the first Seasoar deployment. The associated oxygen sensor was also prone to periods of spiking in the data. For the second deployment, CTD serial number 01-2073 was used, with the exception of the pressure sensor and card, which was removed from CTD 01-1055 and installed in 01-2073. No further sensor problems were reported for the second deployment, with the exception of one recovery of the Seasoar necessitated by biological fouling of the conductivity cell.

#### Miscellaneous

1) The 10kHz pinger used on the 24 way CTD frame was discovered to have a ping repetition rate of greater than once per second (57 pings per minute). This resulted in difficulty in bottom tracking for the deep CTD casts for the first station, but bottom resolution was obtained to the target depth of bottom minus 20 metres for the rest of the casts.

# SHIP'S DAILY LOG

# from Chief Scientist and Master

Date	Time (LT)	Event
01/06/98		Safety Radio Survey completed
02/06/98	0900	17 RVSM staff changeover
		Continue and complete demobilisation
		Commence taking off old trawl wire
03/06/98		Commence mobilisation and taking 271 T of bunkers. Damage to Scintillation
		counter
		Commence winding on new trawl wire
		De-Rat inspection completed
		Trawl wire storage drum shaft sheared
04/06/98		Continue mobilisation and taking bunkers
		IOPP survey completed
05/06/98		Continue mobilisation and taking bunkers
	1500	28 Non-RVSM staff join v/l
	1515	Inspection of galley, fridges and storerooms by North Ayrshire Council (Port
		Health Department)
	1530	Basic Safety Familiarisation briefing for 23 non-RVSM personnel joining in
		Fairlie
	1700	Complete mobilisation
06/06/98		Weather dull and overcast with occasional rain
	0900	Basic Safety Familiarisation briefing for 3 RVS Scientific personnel absent at
		previous day's briefing
		Continue securing
	1136	Clear of berth. No pilot taken
	1212	Clear of Hunterston Channel. Continue securing
		Non-toxic supply turned on mid afternoon
	1600	All secure
	1615-1650	Emergency Drill and Boat Muster
07/06/98	22.50	Weather rougher overnight (F5-6), but moderated during the day
00/07/00	2359	Clocks retarded to GMT
08/06/98	1330	V/I hove to at 59-19.2N 19-00.2W
	1346-1430 1519-1540	CTD CTD
	1625-1704	CTD
	1706	Unable to use Seasoar so proceed towards 59-20N 19-00W
09/06/98	1115	V/I hove to at 59-30N 22-40W, waiting for Seasoar
10/06/98	0000	Reposition to Seasoar start site
10/00/98	0530	V/I hove to at 59-30N 22-30W
	0559-0637	CTD
	0655-0718	Reposition
	0727	Deploy Seasoar in 58-57.9N 22-30.7W. Commence grid survey
11/6/98	0727	Weather warm and sunny, some clouds. Sea pretty calm
12/6/98		Weather somewhat dull but dry, sea pretty flat
13/06/98		Weather sunny all day, the sea really calm
	0557	Seasoar recovered in 59-14.3N 22-22.5W
	0617-1042	Reposition to eddy centre (EC1) at 59-30.3N 21-07.8W
	1047-1142	Test deploy Drifting buoys
	1306-1338	CTD at EC1
	1434	Spar buoy deployed at EC1
	1546	Commence deploying SF6 tracer in 4Nm grid around EC1
	1503	1 <sup>st</sup> Argos buoy deployed
	1606	2 <sup>nd</sup> Argos buoy deployed
	1635	3 <sup>rd</sup> Argos buoy deployed
	1657	4 <sup>th</sup> Argos buoy deployed
14/06/98		Weather fine all day, sea state calm. Several schools of pilot whales sighted close

		to ship
	0043	Complete deployment of SF6 tracer
	0220-0244	CTD at EC1
	0316-0328	CTD at EC1
	0419-0425	CTD at EC1
	0425-1516	Remain hove to at EC1 for air sampling
	1614	Spar buoy recovered due to beacon failure
	1624-	
15/06/98	0226	Underway sampling survey around EC2 now determined at 59-28.7N 21-04.7W
		Weather once again fine and sunny, sea calm.
	0227-0250	CTD at EC2
	0323-0343	CTD at EC2
	0407-0415	CTD at EC2
	0415-1506	Stationary air sampling at EC2
	1505-1526	CTD at EC2
	1602-	
16/06/98	0215	Underway sampling around EC3 now determined at 59-33.9N 21-07.0W
		Weather fine and sunny; sea state calm, becoming oleagenous in the evening
	0216-0235	CTD at EC3
	0305-0320	CTD at EC3
	0401-0407	CTD at EC3
	0415-1300	Stationary air sampling at EC3
•	1344-1440	RIB Workboat away from ship for remote surface sampling
15/07/00	1455-	TI I I I I I I I I I I I I I I I I I I
17/06/98	0155	Underway sampling around EC3
		Weather today somewhat more overcast than recent days, but still warm and with
	0212 0225	occasional sun; sea state still pretty calm
	0213-0235	CTD at EC3
•	0309-0324	CTD at EC3 CTD at EC3
	0416-0422 0428-2110	Stationary air sampling at EC3
	2110-	Stationary an sampling at ECS
18/06/98	0250	Underway sampling around EC4 now determined at 59-33.8N 20-46.1W
		Weather cloudy but bright, sea state a bit rougher than the last few days - slight
		chop. In the evening sea got distinctly rougher
	0302-0321	CTD at EC4
	0353-0355	CTD at EC4
	0411-0424	CTD at EC4
	0436-1542	Stationary air sampling around EC4
	1542-	
19/06/98	0208	Underway sampling around EC4
		Weather and sea state poor all day (F8/9) and planned 48 hour diurnal cycle of
		sampling at patch centre abandoned. Rain collection continues!
	0238-0248	CTD at EC4
	0248	Scientific activities suspended due to adverse conditions. V/l hove to
20/06/98		Gale abated overnight. Weather dull with showers, F4-5.
	1248	Resume scientific activities
	1248-2105	Underway sampling around EC5 now determined at 59-33.3N 21-01.2W
	2210-2232	CTD at EC5
21/06/98		Weather dull all day, sea state moderate
	0101-0302	Underway sampling around EC5
	0302-0320	CTD at EC5
	0348-0359	CTD at EC5
	0359-1506	Stationary air sampling at EC5
	150-1	Eddy centre now determined at 59-34.9N 21-09.3W, EC6
	1507-1533	CTD at EC6
	1611-1614	CTD at EC6
	1642-2140	Underway sampling around EC6. Eddy centre now determined at 59-39N 21-
		20W, EC7

	2140-	
22/06/98	1150	Stationary air sampling at EC7
		Weather rather overcast. Sea calm.
	1150-1158	CTD at EC7
	1200-1223	CTD at EC7
	1223-2014	Reseeding patch with SF6
22/06/09	2015- 0254	Underway sampling around EC7. Eddy centre now determined at 59-54.3N 20-
23/06/98	0234	44.4W, EC8
		Weather dull and rainy with increasing wind, sea lumpy with significant swell.
		Start of 48 hour diurnal.
	0254-0300	CTD at EC8
	0322-0341	CTD at EC8
	0600-0616	CTD at EC8
	0805-0820	CTD at EC8
	1025-1043	CTD at EC8
	1043-1304 1304 <b>-</b> 1306	Survey patch CT at EC8
	1311-1335	CTD at EC8
	1400	Proceed to Vestmannaeyjar to land seaman for compassionate reasons
24/06/98		Wind and sea state moderated during afternoon and evening.
	1300	Off Vestmannaeyjar. Unable to enter due to problem with hand steering control
		Pilot tug collects seaman
	1507	Steering control fixed. Proceed into Vestmannaeyjar entrance
	1518	Waiting 4 cables off harbour entrance for replacement seaman. Unable to land
	1614-1836	scientist to doctor due to adverse conditions
	1836	Missed flight so proceed back out to sea  Proceed into Vestmannaeyjar to collect replacement
	2153	Waiting 3 cables off harbour entrance
	2215-2337	Replacement seaman embarked
	2337	V/l clear of harbour approaches and returning to work area
	2348	
25/06/98	1115-1117	Water sample bottles at 61-27N 20-25W
	1218	After advice from RNH Haslar doctor, decision made to return to
0.640.640.0	0.520	Vestmannaeyjar for scientist to see doctor about eye problem
26/06/98	0530 0710	V/l hove-to off Vestmannaeyjar Scientist disembarked to pilot tug close to harbour entrance
	0923	Scientist disembarked from pilot tug close to harbour entrance
	0936	V/I clear of harbour approaches and returning to work area
	1615-1655	Emergency drill and boat muster
	2106-2128	Water bottle samples at 61-12N 20-31W
27/06/98	0218-0241	CTD at 60-26N 20-40W
	0308-0310	Water sample bottles at 60-26N 20-40W
	0326-0345	CTD at 60-26N 20-40W
	0422	Resume passage to work area
	0907-2120	SF6 patch survey CTD at 59-45.7N 20-59.5W
	2129-2148 2226-	C1D at 59-45./N 20-59.5 W
28/06/98	0152	SF6 patch survey
20,00,00	****	Weather generally fine but somewhat overcast, with a rain shower in the
		afternoon. Sea state pretty calm.
	0208-0233	CTD at 59-48.4N 20-58.0W
	0319-0341	CTD at 59-48.4N 20-58.0W
	0423	Commence drifting buoy search
	0905	Recover RVS 1954 buoy
	0933 1002	Recover RVS 1953 buoy Recover DML buoy. Resume search for RVS 03945 buoy
	1405	Cease search. DF USELESS
	1418-1501	CTD at 59-50.0N 20-41.0W

	1506-1508 1557	Seasoar deployed. Commence grid survey
29/06/98		Sea calm, weather overcast.
		Continue Seasoar grid survey
30/6/98		Weather warm and calm, with sun in the evening. Sea state calm.
		Continue Seasoar grid survey
01/07/98		Sea calm, weather overcast but only light wind and quite warm.
	0740	Seasoar recovered due to dirty sensor
		Seasoar redeployed. Resume grid survey
02/07/98		Weather quite windy (F5), sea moderately choppy.
	0855	Seasoar recovered. Commence search for missing Argos buoy 03945
	1308	Search for buoy cancelled due to poor Argos position information and poor DF
	1010 1670	determination signals
	1310-1652	Reposition to start of CTD transect across eddy
	1704-1709	CTD at 59-59.8N 20-03.8W
	1710-1730	Reposition
	1736-1940	CTD at 60-00.0N 20-04.3W
	1949-2122	Reposition CTD at 59-53.2N 20-21,6W
	2125-2325 2345	CTD at 59-53.2N 20-21.6 W
03/07/98	0115	Reposition
		Weather similar to 2/7/98
	0121-0306	CTD at 59-46.4N 20-39.2W
	0310-0439	Reposition
	0443-0634	CTD at 59.39.7N 20-57.0W
	0641-0805	Reposition
	0815-1013	CTD at 59-32.5N 21-13.9W
	1020-1130	Reposition
	1140-1336	CTD at 59-25.7N 21-31.4W
	1340-1450	Reposition
	1453-1646	CTD at 59-18.8N 21-48.4W
	1650-1818	Reposition
	1822-2014	CTD at 59-11.6N 22-05.4W
	2018-2214	Reposition
04/07/00	2218-	OTD 4 60 04 51 00 01 011
04/07/98	0008	CTD at 59-04.5N 22-21.9W
	0008-0400	High productivity survey
	0442-0453 0512-0533	CTD at 59-20.4N 21-44.6W CTD at 59-20.3N 21-44.5W
	0533-1830	Stationary air sampling
	1834-1841	CTD at 59-20.4N 21-55.7W
	1900	Proceed towards Southampton
05/07/98	1500	Weather overcast but calm and reasonably warm
00,07190	0654-0712	Water bottle at 57-39.2N 19-24.2W
06/07/98		Weather calm but overcast
	0612-0833	CTD at 54-24.6N 14-50.2W
	0841-0843	Water bottle at 54-24.2N 14-50.3W
	1615-1640	Emergency drill and boat muster
	1900-1912	Water bottle at 52-58.6N 12-57.8W
07/07/98		Weather calm but overcast.
	0650-0702	Water bottle at 51.27.9N 10-52.9W
	1848-1906	Water bottle at 50-16.9N 7-56.0W
08/08/98		Weather warm and sunny.
	1830	Entering Needles Channel
	1952	Pilot on board at East Lepe
	2106	V/l secure stbd side to SOC berth, Empress Dock

# **SAMPLES COLLECTED DURING D234**

Tables compiled by Polly Machin and Wendy Broadgate

- Table 1 Underway Water Sampling Log
- Table 2 Log of Events CTD, Hydrowire, Underway, Air, XBTs
- Table 3 CTD Stations Date, Lat, Long, Depths

						ierway v	va.co						
Sample	Date/ Time (GMT)	chl/pigs	nuts	hydro	halo	N20/CH4	bac	phyto	DMS	sal	Taken by?	lat	lon
PG001	09/06/98 09:31					у						59.49058	-22.30255
PG002	09/06/98 09:37				.,	у						59.49159	-22.32546
PG003	09/06/98 10:11					у			v			59.49581	-22.44829
PG004	09/06/98 10:14					у						59.49619	-22.45903
PG005	09/06/98 11:12				.,	у						59.50593	-22.66487
PG006	09/06/98 12:56					у		.,,,,,,,,,				59.51863	-22.66508
PG007	09/06/98 14:03					у						59.52162	-22.64683
PG008	10/06/98 06:35							AFC				58.99236	-22.50809
PG009	10/06/98 06:37	11 (chl)		\$5.02	у							58.99217	-22.50853
PG010	10/06/98 09:00	12 (chl)	12		*******					145		58.99406	-22.54231
PG011	10/06/98 11:00	13 (chl)				4.044.4		AFC2				59.00296	-22.09881
PG012	10/06/98 11:13		13							146		59.00341	-22.04980
PG013	10/06/98 11:18							У				59.00336	-22.03096
PG014	10/06/98 13:00	14 (chl)	14			***********				147	<u> </u>	58.99992	-21.64128
PG015	10/06/98 13:05							AFC3				58.99983	-21.62208
PG016	10/06/98 15:00	15 (chl)										59.00051	-21.18720
PG017	10/06/98 15:01		15					AFC4				59.00053	-21.18353
PG018								<u> </u>					
PG019	10/06/98 15:03									148		59.00051	-21.17612
PG020	10/06/98 17:03	16 (chl)	16	\$4.03	у			y AFC5		149		58.99819	-20.73081
PG021	10/06/98 19:00	17 (chl)	17	S1.03	У			y AFC6	S1	150		59.10005	-20,49884
PG022	10/06/98 21:00	18 (chl)	18	\$2.03	у		<u></u>	y AFC7	S2	151		59,19947	-20.76832
PG023	10/06/98 22:59	19 (chl)	19	\$2.04	у			AFC8	S3	152		59.20278	-21.26619
PG024	11/06/98 01:00	20	20						S4	153		59.20125	-21.76887
PG025	11/06/98 03:00	21	21							154		59.20042	-22.26237
PG026	11/06/98 05:00	22 (chl)	22							155		59.32789	-22.49708
PG027	11/06/98 07:00	23 (chl)	23					AFC9		156		59.40092	-22.15385
PG028	11/06/98 08:59	24 (chl)	24	\$2.05	у			AFC10	S5	157		59.40534	-21.69618
PG029	11/06/98 09:06					у						59,40535	-21.66979
PG030	11/06/98 09:09					у						59.40535	-21.65856
PG031	11/06/98 10:58	25 (chl)	25	S2.06	,	у		AFC11	S6	158		59.39853	-21,24949
PG032	11/06/98 12:51		26									59.39678	-20.83466
PG033	11/06/98 12:59	26 (chl	)	\$2.07	у			AFC12	\$7	159		59.39699	-20.80503
PG034	11/06/98 14:58	27	27	\$2.08	у			AFC13	S8	160		59.47377	-20.49790
PG035			28	\$2.09	у			AFC14	S9			59.59911	-20.69568
PG036		28								161		59.59924	
PG037	1115555 1000	29	1	\$2.10	) у				S10	162		59.60089	<del></del>
PG038		1	1					AFC15				59.60084	
PG039		30	2		у			AFC16		163		59.60327	-21.56073
PG040		31	3		у			AFC17		165	sal bottle broke	59.60138	-21.98865
PG041		<del></del>	4							166		59,59922	-22.42539
PG042		<del></del>	5							167		59.80010	-22.46628
PG043		34	6							168		59.80177	-21.95815
PG04			7		1			AFC8	S11	97		59.80127	-21.45067
PG045						UW011					tom	59.79798	-20.97986
PG046			8	S2.1	Ĭ	UW012		AFC9	S12	98		59.79794	
PG04			1	1		UW013						59.79700	
PG048			1	1		UW014	1					59.79580	
PG04			1	1		UW015						59.79547	-20.51719
PG050			K 9	S2.1	2 y	UW016		AFC20	S13	99	DMSP and lyase	59.83271	-20.49755
PG05			1		T	UW017						59.89257	-20.49780
PG05			1		1	UW018	"					60.00075	
PG05			10	\$2.1	3 у	UW019		AFC21	S14	100		60.00029	-20.64658
PG05					1								
PG05		1	<b></b>	<b>—</b>	1	UW020						60,00019	-20,75913
PG05		<del></del>	-			UW021						60.00142	-20.98559
PG05			11	\$2.1	4 y							60.00187	-21.08935
PG05			+ ''	<del></del>	<b>-</b>				S15	5		60.00155	-21.11854
PG05			12	S2.1	5 y	UW023		AFC23		<del></del>		59.99860	
PG06			12	- JZ. 1	- ,	UW024			1			59.99788	-21.63999
PG06	····				+	UW025						59.99966	-21.80868
FOU	1 12/00/80 10,30	<u>,                                    </u>				1 011020		,					

·····		·		<del></del>	·····	Jeiway v	VUIC-	ı Samp	7111113	109	· · · · · · · · · · · · · · · · · · ·		
Sample	Date/ Time (GMT)	chl/pigs	nuts	hydro	halo	N20/CH4	bac	phyto	DMS	sal	Taken by?	lat	lon
PG062	12/06/98 18:57		13							103		60.00016	-21.90322
PG063	12/06/98 19:03	42		S2.16	у	UW026		AFC24	S17			60.00019	-21.92484
PG064	12/06/98 19:40					UW027						59.96847	-22.01004
PG065	12/06/98 20:04					UW028						59.92572	-22.03242
PG066	12/06/98 21:01	44	14	S2.17	У	UW029		AFC25	S19	104	ļ	59.83681	-21.96453
PG067	12/06/98 23:00	45	15							105		59.68751	-21.57491
PG068	13/06/98 01:00	46	16							106		59.53621	-21.20706
PG069	13/06/98 03:00	47	17							107		59,38864	-20.85245
1												ļ	
PG070	13/06/98 05:05	48	18							108		59,25118	-20.50245
PG071	13/06/98 05:54									109	SeaSoar 5m	59.23828	-20.37664
PG072	13/06/98 07:15							AFC26				59.25500	-20.53216
PG073	13/06/98 09:40					UW030	<u></u>					59.39378	-20.91230
PG074	13/06/98 10:05					UW031						59.43614	-21.00289
PG075	13/06/98 11:32					UW032						59.51122	-21.13514
PG076													
PG077	14/06/98 16:03	71	11	\$3.03		UW033			S20			59.47853	~21.06055
PG078	14/06/98 16:37					UW034						59.45913	-21.05089
PG079	14/06/98 17:03	72	12	\$3.04	V	UW035		1	S21			59,42995	-21.02873
PG080	14/06/98 17:35	7 844	1 / /	00.01		UW036			<del> </del>			59.43022	-21.18532
PG081	14/06/98 18:05	73	13	\$3.05		UW037			\$22			59.45542	-21.26329
	£	13	13	33.00					322	~		<del></del>	
PG082	14/06/98 18:35					UW038			600			59.45496	-21.13746
PG083	14/06/98 19:02	82	14	\$3.06		UW039			S23			59.45434	-21.02424
PG084	14/06/98 19:32					UW040	ļ	2	ļ			59.48073	-21.06751
PG085	14/06/98 20:07	83	15	S3.07		UW041			S24			59.48154	-21.21085
PG086	14/06/98 20:38	<u> </u>		L		UW042						59.50624	-21.24936
PG087	14/06/98 21:00	84	16	\$3.08		UW043		3	S25			59.50550	-21.16523
PG088	14/06/98 21:30					UW044						59.50088	-21.05155
PG089	14/06/98 22:01	85	17	\$3.09	у	UW045			\$26			59.53037	-21.03541
PG090	14/06/98 22:31					UW046						59.53056	-21.14853
PG091	14/06/98 23:00	86	18	\$3.10	У	UW047		4	S27			59.55678	-21,10056
PG092	14/06/98 23:30				<b>'</b>	UW048		-/-				59.50987	-21.07320
PG093	14/06/98 23:59	87	19	\$3,11	у	UW049			S28			59.45190	-21.07209
PG094	15/06/98 00:30		10	00.11	,	UW050			020			59.40302	-21.05228
		00						5	S29			59.40447	-20.94631
PG095	15/06/98 01:00	88	20	ļ	У	UW051	د.،	3	329				£
PG096	15/06/98 01:25				-	UW052	-		ļ			59.44670	-20.94260
PG097	15/06/98 02:23	89	21	ļ			ļ					59.47900	-21.08038
PG098	15/06/98 16:00	103	1							****************		59,55435	<del></del>
PG099	15/06/98 17:05	106	2	\$1.06		UW053		6	\$30			59.55600	-21.20189
PG100	15/06/98 18:01	107	3			UW054			S31			59.57965	-21.16376
PG101	15/06/98 19:02	108	4			UW055		7	S32			59.54583	-20.97824
PG102	15/06/98 20:06	109	5	\$2.20	У	UW056	ļ		833			59.53156	-21.20439
PG103	15/06/98 21:01	110	6	1	y	UW057		`8	S34			59.50733	-21.18862
PG104	<del> </del>			<u> </u>	-	UW058	<b>†</b>					59.50484	-21.07009
PG105		111	7	\$5.05	у	UW059	<del>                                     </del>		S35			59,49597	-20.97734
PG106	15/06/98 23:00	112	8	100.00	+	UW060	<del> </del>	9	S36			59.51244	-21.13479
PG107		112		ļ	У	UW061		K	330			59.57544	-21.13397
<u> </u>	15/06/98 23:30	140	<del></del>		1	<del> </del>	<del> </del>		027				
PG108	16/06/98 00:01	113	9	ļ	У	UW062	-	<del> </del>	S37	-	-	59.60483	-21.06181
PG109	16/06/98 00:30		ļ	ļ	ļ	UW063	ļ		\$38			59.62007	-20.97659
PG110	<del></del>	114	10	S5.06	У	UW064		10	S39	ļ		59.63053	-21.07472
PG111	16/06/98 01:30		<u> </u>		ļ	UW065		-	S40			59.59744	-21.15366
PG112	16/06/98 02:00	115	11		у	UW066			\$41			59.56541	-21.11941
PG113	16/06/98 16:00	128	1	\$2.22	у	UW067			\$42			59.60711	-20.92318
PG114	16/06/98 16:30					flask smashe	d		\$43			59.60825	-20.98299
PG115	16/06/98 17:00	129	2	\$2.23	у	UW068		14	S44			59,62940	-21.05568
PG116	£	130	3	\$1.08	- <del> </del>	UW069	T		\$45			59.63219	-20.80603
PG117	<u> </u>	131	<u> </u>		<del> </del>	UW070	1		\$46			59.59347	-20.73957
PG118	1	132	4		у	UW071	+	.15	\$47		1	59.58391	-20.82058
PG119	<del></del>	133			<u> </u>	·	-	1 13				59.58391	-20.93570
	<del>                                     </del>		<del>  </del>	04.00	-	UW072	-		\$48		<del></del>	<del></del>	
PG120	16/06/98 20:00	134	5	S1.09	У	UW073	<del> </del>	ļ	\$49	-		59.58450	-21.05185
PG121	16/06/98 20:30	141		1	I	UW074 UW075	ļ		S50 S51	<u> </u>		59.55678 59.55671	-21.03562
PG122	16/06/98 21:00	142	6	S2.24	у			16					-20.91735

						ierway w						1-4	1
	Date/ Time (GMT)	chl/pigs	nuts	hydro	halo		bac	phyto	DMS	sal	Taken by?	lat	lon
PG123	16/06/98 21:30	143				UW076			\$52			59.55502	-20.79775
PG124	16/06/98 22:00	144	7	S1.10	у	UW077			S53			59.53709	-20.71021
PG125	16/06/98 22:31	145				UW078			S54			59.53219	-20.82659
PG126	16/06/98 23:03	146	8			UW079		17	S55			59.53325	-20.9561Ò
PG127	16/06/98 23:30	147				UW080			S56			59.50739	-20.94694
PG128	17/06/98 00:01	148	9			UW081			857			59.50577	-20.82505
PG129	17/06/98 00:30	149				UW082			\$58			59.50468	-20.71103
PG130	17/06/98 01:01	150	10			UW083		18	S59			59,53751	-20.65933
PG131	17/06/98 01:35	151				UW084			\$60			59.56248	-20,72834
PG132	17/06/98 01:59	152	11			UW085			S61			59.56384	-20.76833
PG133	17/06/98 21:10	164	1		У	UW086			\$62			59.46775	-20.78956
PG134	17/06/98 21:30	165				UW087			S63			59.43994	-20.80093
PG135	17/06/98 22:00	166	2		У	UW088			\$64			59.43343	-20.88315
PG136	17/06/98 22:30	167			,	UW089			S65			59.46716	-20.90434
PG137	17/06/98 23:00	168	3	s2.25	у	UW090		19	\$66	·		59.47962	-20.86320
PG138	17/06/98 23:30	169	<u> </u>	52.20		UW091			S67			59,43462	-20,84713
<u> </u>	17/06/98 00:01	170	4		У	UW092	-		-			59.50577	-20.82505
PG139		171	-		у	UW093	1		S68		-	59.43310	-21.10324
PG140		·	5	-1 11		UW094	ļ	20	\$69			59.50145	-21.09826
PG141	18/06/98 01:00	172	5	s1.11				20	S70			59.50019	-20.98134
PG142	18/06/98 01:30	173		00.00		UW095			310			59.50011	-20.96560
PG143			ļ	\$2.26	/ <del></del>	1011000			0.74			59.50411	-20.86867
PG144	<del></del>	174	-	S3.15		UW096		-	\$71			59.53411	-20.95360
PG145		175		ļ		UW097			\$72		<u> </u>		
PG146	18/06/98 16:03	186	1	\$4.08	У	UW098	ļ		873			59.55688	-20.93999
PG147	18/06/98 16:31					UW099						59,58099	-20,96734
PG148	18/06/98 17:00	187	2		у	UW100		22	S74			59.59113	-20.92041
PG149	18/06/98 17:30	188			ļ	UW101			\$75			59.56603	-20.89972
PG150	18/06/98 18:34	189			у	UW102			S76			59,58991	-20.99672
PG151	18/06/98 19:00	190	3	\$4.09		UW103		23	S77			59.60798	-20.93508
PG152	18/06/98 19:20				у							59.60526	-20.86578
PG153	18/06/98 19:32	191				UW104			S78			59.58418	-20.86646
PG154	18/06/98 20:02	192	4	\$4.10		UW105			579			59.52516	-20.87121
PG155		193				UW106			\$80			59.52629	-21.01185
PG156		194	5	S1.12		UW107		24	S81			59.58102	-21.03876
PG157		195				UW108			\$82			59.62492	-20.99502
PG158		196		S4.11	ļ	UW109			\$83			59,62609	-20.94962
PG159		197				UW110			S84			59.64379	-20.95782
PG160		198	6	\$4.12		UW111	1	25	S85			59.64476	-21.08132
PG161		199	+- <u>`</u> -	J 1.12	ļ	UW112	-	<del> </del>	S86			59.63601	-21.17663
PG162		200	+7	\$4,13		UW113	<del> </del>	<del></del>	S87			59.60272	-21.17136
		201	+ '-	04.10	-	- 000110	+	-	588			59.60280	
PG163			8	-	┼	UW114	<del></del>	26	S89		<u> </u>	59.66359	
PG164		202			-	000114			- 003			59,68564	
PG16		-		-	У		-	-	\$90	+		59.67158	
PG166		<del></del>	9	-	у	UW115	-	-	780	+		59,61963	
PG16					У	1.58/4/4						59.55126	
PG168						UW116	-		-			59.551242	
PG169					У	<del> </del>		-	-				
PG170					ļ	UW117						59.59733	
PG17			11	\$1.13	+	UW118	4	28	S91	<u> </u>		59.60178	
PG17	2 20/06/98 18:00	207	12	<del></del>	<del></del>	UW119		-	S92			59.56047	
PG17	3 20/06/98 19:00	208	13	\$4.14	у			29	S93			59.55485	
PG17	4 20/06/98 19:32	209				UW121			S94			59.52926	
PG17	5 20/06/98 20:00	210	14		У		<u> </u>			1		59.52723	
PG17	6 20/06/98 20:03			\$2.28	3	UW122			\$95			59.52715	
PG17	7 20/06/98 20:30	211				UW123			\$96			59,52732	
PG17	8 20/06/98 21:00	212	15	\$4.15	у			30	\$97			59.52949	
PG17	9 20/06/98 22:28	214	17	\$2.29	у	UW124			598			59.54857	
PG18			10		+	UW125		42	S <b>9</b> 9			59.63003	-21.41667
PG18					1	UW126			\$100	)		59.62141	-21.30354
PG18			11	83.19	) y		1		\$10			59.59828	3 -21.38061
PG18					y	4111400			\$102	<del></del>		59.60679	-21.51025
L- 310	5 2 1700/30 ZU.SU	<u> </u>			1 9	1 077720							

				<i>7</i> 207	Oil	Jervay	valo	Jann	mig	109		·	
Sample	Date/ Time (GMT)	chl/pigs	nuts	hydro	halo	N20/CH4	bac	phyto	DMS	sal	Taken by?	lat	lon
PG184	21/06/98 21:00	у	12		у	UW129		47	S103			59.65246	-21.46979
PG185	21/06/98 21:30	у			у	UW130			\$104		,	59.65195	-21.34081
PG186	22/06/98 14:15					UW131						59.91260	-21.16943
PG187	22/06/98 15:15		************			UW132						59.93777	-21.16712
PG188	22/06/98 16:00					UW133	<del> </del>					59.91483	-21.14630
PG189	22/06/98 17:00	-				UW134						59,90693	-21.06322
PG190	22/06/98 18:15					UW135						59.93914	-20.99551
PG191	22/06/98 18:44					UW136						59.95582	-20.93331
	,	057	4		ļ				0105		***************************************	<del></del>	
PG192	22/06/98 19:00	257	1		у	UW137			S105			59.96326	-20.97404
PG193	22/06/98 19:36	258				UW138	~		S106			59.96163	-20.99771
PG194	22/06/98 20:01	259			У	UW139			S107			59.96003	-21.01374
PG195	22/06/98 20:35	260			У	UW140			S108			59,95839	-21.03631
PG196	22/06/98 21:02	261	2			UW141		56	\$109			59.95732	-21.05553
PG197	22/06/98 21:30	262	ļ						S110			59,95663	<i>-</i> 21,07486
PG198	22/06/98 22:00	263							S111	-		59.95585	-21.00281
PG199	22/06/98 22:30	264							S112			59.91632	-20.96078
PG200	22/06/98 23:02	265	3					57	S113			59,91818	-20.89681
PG201	22/06/98 23:30	266							S114			59.95053	-20.81384
PG202	23/06/98 00:01	267					1		\$115			59.92346	-20.77318
PG203	23/06/98 00:31	268				<del> </del>	+	l	S116		<del></del>	59.90706	-20.79963
PG204	23/06/98 01:00	269	4					58	S117			59.94399	-20.86312
PG205	23/06/98 01:31	270			***************************************				\$118			59,93995	-20.86441
	· · · · · · · · · · · · · · · · · · ·	<del>                                     </del>	<del></del>				<del> </del>					· <del> </del>	-20.83949
PG206	23/06/98 02:00	271				11144440			S119			59.91704	
PG207	23/06/98 02:32	<u>y</u>				UW142			S120			59.90814	-20.77762
PG208	23/06/98 03:35	У			У							59.90521	-20.73821
PG209	25/06/98 00:01	305	17	\$2.37	у	UW143		84	S121		lyase	63.41313	-20.13305
PG210	25/06/98 00:21	308				UW144			\$122			63,34961	-20.13785
PG211	25/06/98 00:41	306			У	UW145		AFC	S123		flow ch 2	63.28443	-20.14866
PG212	25/06/98 01:00	307	18	S2.48	у	UW146		85/AFC	S124		lyase	63.22382	-20.15723
PG213	25/06/98 01:20	309				UW147		AFC	\$125			63,16084	-20.16666
PG214	25/06/98 01:39	310			у	UW148		AFC	S126		***************************************	63.10060	-20.17504
PG215	25/06/98 02:00	311	19	\$2.39	у	UW149		86/AFC	S127		lyase	63.03520	-20.18420
PG216	25/06/98 02:16	312			<u> </u>	UW150	1	<u> </u>	S128		······································	62.98537	-20.19670
PG217	25/06/98 02:30	313			У	UW151		AFC	S129			62.94184	-20.20677
PG218	25/06/98 02:45	314			<b>–</b>	UW152			S130			62.89517	-20.21321
PG219	25/06/98 03:00	315	20	\$2.40	у	UW153		87/AFC	\$131			62.84886	-20.21884
PG220	25/06/98 14:00	317	22	\$2.42		011100		89/AFC				61.55928	
PG221	25/06/98 15:04	318	23	S2.43		UW154	+	90/AFC	·			61.72004	-20.38848
PG222		<del></del>	_20	32.40		UW155		SUMPC	\$134		<del></del>	61.80113	-20.38064
	25/06/98 15:35	320				·						<del>-</del>	
PG223	25/06/98 16:34	322				UW156			\$135			61.86854	-20.36940
PG224	<del> </del>	323	25	ļ	<u> </u>	UW157	<u> </u>	92	S136		CO2	61.93488	-20.35472
PG225	25/06/98 18:00	324	26	S2.45	У	UW158		93/AFC	\$137			62,09394	-20.32449
PG226	25/06/98 18:30	326		ļ	<u> </u>	UW159			S138			62.17418	-20.30416
PG227	25/06/98 19:31	327	<u> </u>		ļ	UW160			S139			62.25034	-20.26446
PG228	25/06/98 20:01	328		\$2.46		UW161			S140			62.31704	-20.24338
PG229	25/06/98 20:04		28					95/AFC				62.32328	-20.24142
PG230	25/06/98 20:58	329	1	\$4.22	у	UW162		96/AFC	\$141			62.45287	-20.23056
PG231	25/06/98 21:32	330	1		у	UW163	T		\$142			62.53824	-20.24818
PG232	25/06/98 22:32	333			у	UW164	1		S143		·····	62.61113	-20.24046
PG233	25/06/98 23:00	332	3	S2.48	<del></del>	UW165	+	98	S144			62.67951	-20.23394
PG234	<del></del>	334	<del>                                     </del>	J2.70	y	UW166		<del>  ~~</del>	S145			62.75354	-20.21851
PG235		225	4	\$4.23	+	UW167	-	99	\$146	-		62.83288	-20.20635
PG236		220		04.23	у	OVV 10/		22	3140	2		61.36990	<del></del>
	<del> </del>	-		00.45	<del>  -</del>	108/400	-	-	04.17	L			-20.50204
PG237	26/06/98 21:25		ļ	S2.49	У	UW168			S147			61.19748	-20.51596
PG238	27/06/98 00:02	<u> </u>				ļ			<b></b>	3		60.83126	
PG239	27/06/98 03:59			<u> </u>		]	<u> </u>		<u> </u>	6		60.42691	-20.66558
PG240	27/06/98 23:02	359		\$1.24				139	S148			59.80339	-21.06989
							1	1	10440			FO 04 400	-20.86216
PG241	28/06/98 00:08	360		\$1.25				140	S149			59.81433	-20.00270
		360 361		\$1.25 \$1.26			+	140	\$149			59.81433	-20.96655
PG241	28/06/98 00:08 28/06/98 01:30	+						<del> </del>	+				

			1	JZ-04	One	ucivay v	Valor	Samp	11119	<del>- 09</del>		,	
Sample [	Date/ Time (GMT	chl/pigs	nuts	hydro	halo	N20/CH4	bac	phyto	DMS	sal	Taken by?	lat	lon
PG245	28/06/98 17:04		27							60		59.91937	-20.68222
PG246	28/06/98 18:04		28									59.98153	-20.84644
PG247	28/06/98 19:00		1							61		60.04080	-21.01555
PG248	28/06/98 20:00		2							62		60.10451	-21.20001
PG249	28/06/98 20:59		3							63		60.16313	-21.38975
PG250	28/06/98 22:00		4		<b></b>		<b></b>	-		64		60.14451	-21.50038
		-	5		<del> </del>		-			65		60.03850	-21.49848
PG251	28/06/98 23:00		6		<del> </del>					66	ļ	59.92431	-21.49879
PG252	29/06/98 00:00	-			-					67		59.81584	-21.49967
PG253	29/06/98 01:00		7	ļ	ļ		+-+					59.70713	-21.50144
PG254	29/06/98 02:00		8		ļ		<u> </u>		-	68	-		-21.50153
PG255	29/06/98 03:00		9		ļ		4			69	ļ. <u></u>	59.60096	
PG256	29/06/98 04:00		10				ļ			70		59.49726	-21.49978
PG257	29/06/98 05:00		11							71		59.39051	-21.49747
PG258	29/06/98 06:00		12							72		59.28070	-21.50051
PG259	29/06/98 07:00		13							73		59.19989	-21.43544
PG260	29/06/98 08:03		14							74		59.24687	-21.30135
PG261	29/06/98 09:00	-	15							75		59.36159	-21.30132
PG262	29/06/98 10:00		16	1	1					76		59,50000	-21.30797
PG263	29/06/98 11:00	+	17	<del>                                     </del>	<del> </del>	-	1			77		59.63850	-21.30176
PG264	29/06/98 12:03	-	29		-	<del> </del>			<del> </del>	78		59.78076	-21.30095
			18			-	45		<del> </del>	79		59.90708	-21.30002
PG265	29/06/98 13:00	<del></del>		<del>-</del>	<del> </del>	<b>_</b>	46		<del> </del>	80		60,03805	-21.29996
PG266	29/06/98 14:00		19		-		4		<del> </del>	81	-	60.16072	-21.29976
PG267	29/06/98 15:00		20		ــــــ		47	450		01		60.16694	-21.29988
PG268	29/06/98 15:03	_ <del></del>				UW169		156	S31				
PG269	29/06/98 15:59	386				UW170	48	157	\$152			60.20002	-21.12604
PG270	29/06/98 16:02		21	S5.17				ĺ		82	ļ	60.19919	-21.11334
PG271	29/06/98 16:58	387		\$2.52	2	UW171	49	158	\$153			60.09021	-21.10004
PG272	29/06/98 17:00		22							83		60.08633	-21.10016
PG273	29/06/98 18:00		23						ļ <u>.</u>	84		59.97419	-21.09742
PG274	29/06/98 18:35					UW172						59.91251	-21.09949
PG275	29/06/98 18:38		1		-				S154			59.90727	-21.09976
PG276	29/06/98 19:01		24	\$2.53	3	UW173	50	159	S155	85		59,86716	-21.10020
PG277	29/06/98 20:01		25	\$2.54		UW174		160	\$156			59.75732	-21.10154
	29/06/98 21:02		<del> </del>	\$2.55		UW175		161	S157			59,63543	-21.10178
PG278			у 26	32.3		000173	<del> </del>	101	10,07	12		59.63910	
PG279	29/06/98 21:00						-	<del> </del>	+	87		59.52164	
PG280			27	_				ļ		88		59.40172	<del></del>
PG281			28		-			<del> </del>				59.92431	
PG282	30/06/98 00:00	<i>!</i>	1					ļ	<u> </u>	89			· · · · · · · · · · · · · · · · · · ·
PG283	30/06/98 01:00	)	2					ļ		90		59.20033	
PG284	30/06/98 02:00	)	3					ļ		91		59.28027	
PG285	30/06/98 03:00	)	4							92		59,42640	
PG286	30/06/98 04:00	,	5							93		59,57681	
PG287			6				T			94		59.72790	
PG288			7	1		1				95		59.87719	-20.90026
PG289			8	-					1	96		60.01864	-20.89915
PG290			9				1		1	13		60.15348	-20.89706
			10		+		_	<del> </del>	S158			60.28368	-20,90330
PG291			- -10		-		<del></del>		10.00	15		60,38393	
PG292			+	-	-			+	\$159			60.41082	
PG293			11	-			+	-				60,49669	
PG294			12		<del> </del>		S1		S160			60.40188	
PG295	30/06/98 12:00	0 4/397	7 13				S2		\$161	<del></del>			
PG296	30/06/98 13:00	5/394	4	S2.5	6	UW176	S3	162	\$162			60.26117	
PG297	7 30/06/98 13:00	3	14							19		60.25411	
PG298	3 30/06/98 14:00	0 1/395	5 15	S2.5	.7	UW177	\$4	163	\$163	3 20		60.11920	
PG299		<del></del>	6 16	S2.5	8	UW178	\$5	164	S164	4 21		59.98054	4 -20.69943
PG300						UW179	<del></del>	AFC/16	55 S165	5 22		59.84803	-20.70090
PG301				+ 55.2	+-	UW180		1		1		59.78429	9 -20.69947
			0 40	637	90	UW181		AEC/46	36 S166	6 23		59.72192	
PG302	<del></del>								57 S16			59.60129	
PG303				S3.2	. <del>a</del>	UW182		AFU/10	11 310			59.59916	
PG304	4 30/06/98 18:0	0	19	)						24		59,4749	
PG305		0 401	20	S3.3		UW183		AFC/16		8 97			

Sample	Date/ Time (GMT)	chl/pigs				N20/CH4		<del>,                                    </del>		sal	Taken by?	lat	lon
PG306	30/06/98 20:00	-impige	21	.ry at O	naio	.120,0114	- ac	Piijto	٥.,١٥	98	i uncil by :	59.34615	-20.70618
PG307	30/06/98 20:09	402	<u></u>									59.32666	-20.70422
PG308	30/06/98 21:00	702	22							115		59.21457	-20.70208
PG309	30/06/98 22:00		23				ļ			99		59.21453	-20.70200
PG310	30/06/98 23:00		24				1			100		59.35572	-20.50226
PG311	01/07/98 00:00		25				-			101	-	59.27874	-21.10228
PG311	01/07/98 01:00		26							102		59.64483	-20.49825
PG312	01/07/98 02:00		27				-			103		59.78746	-20.49835
PG314	01/07/98 03:00		28							103		59.92488	-20,49033
PG315	01/07/98 04:00		1							105		60.05884	-20.49892
PG316	01/07/98 05:00		2			~~~~~				106		60.19160	-20.50091
PG317A	01/07/98 06:00	-	3				<del> </del>			107		60.31779	-20.30091
PG317	01/07/98 09:05			-,			<del> </del> -			107	SeaSoar 5m	60.15327	-20.49943 -20.53629
PG318	01/07/98 10:00		4							109	Seasoai Sili	60.13327	-20.50252
PG319	01/07/98 11:00		5				-			110	<u> </u>	60.25459	-20.30232 -20.49861
PG320	***************************************		 6				-			111		60.36677	
	01/07/98 11:50						-						-20.49657
PG321	01/07/98 13:01 01/07/98 14:00	-	7 8					ļi		112		60.49908	-20.43787
PG322	F TOTAL CO. \$4000 ALL FOR \$500 ALL FAMILY \$440 ALL FOR \$500 ALL FOR \$5		9				<b></b>			113		60.42481	-20.29954
PG323	01/07/98 15:00	40/444	9	00.04				100		116		60.27935	-20.30108
PG324	01/07/98 15:06	10/411		S3.31			S6	169				60.26486	-20.30004
PG325	01/07/98 15:42	412	-10				ļ			447		60.17831	-20.29990
PG326	01/07/98 16:00	407	10	00.00			ļ	470/450/	0400	117		60.13469	-20.29995
PG327	01/07/98 16:59	427		\$3.32			ļ	170/AFC1	\$169	440		59.99104	-20.29875
PG328	01/07/98 17:00	440	11	00.50	ļ	104404	ļ	1744506	0470	118	<del> </del>	59.98868	-20.29883
PG329	01/07/98 18:04	443	12	\$2.59	+	UW184	-	171/AFC2		119		59.83501	-20.30023
PG330	01/07/98 19:00	444	13	\$3.33		UW185	57	172/AFC	F	120		59.70150	-20,30204
PG331	01/07/98 19:20					UW187		4501.5				59.65538	-20.30021
PG332	01/07/98 20:02		14	S2.60		UW187	58	173/afc	S171	49		59.55849	-20.30088
PG333	01/07/98 20:59						59					59.42902	-20.30098
PG334	01/07/98 21:01		15			UW188	ļ	174/afc	S172	50		59.42454	-20.30135
PG335	01/07/98 21:10			S3.34	I		ļ					59.40464	-20.30312
PG336	01/07/98 21:46			\$4.28	·		1		S173			59,32304	-20.30428
PG337	01/07/98 21:50	ļ		\$1.28			ļ		S174			59.31418	-20.30402
PG338	01/07/98 22:00		16	<u> </u>				<u> </u>		51		59.29179	-20.30295
PG339	01/07/98 22:02			ļ				afc				59.28732	-20,30266
PG340	01/07/98 23:00		17		ļ		ļ			52		59.20030	-20.20594
PG341	01/07/98 23:17			ļ	<u> </u>				ļ			59,19955	
PG342	02/07/98 00:00		18					<u> </u>		53		59.28650	-20.10027
PG343	02/07/98 01:00		19							54		59.41966	-20.10040
PG344	02/07/98 02:00		20			~~				55		59.54931	-20.09865
PG345	02/07/98 03:00		21		ļ		<u> </u>			57		59.66996	-20.09828
PG346	02/07/98 04:00		22							58		59,78694	-20.09958
PG347	02/07/98 05:00	ļ	23		<b> </b>					59		59,90281	-20.09992
PG348	02/07/98 05:14	473			<b> </b>		1		S175			59,89757	-20.12780
PG349	02/07/98 05:15			ļ	ļ				S176			59.89547	-20.12780
PG350	02/07/98 05:41	474			<b></b>		ļ		S177			59.83824	-20.12136
PG351	02/07/98 05:42	<u> </u>		ļ	<u> </u>				S178			59.83612	-20,12128
PG352	02/07/98 06:00		24							60		59.79841	-20.12988
PG353	02/07/98 06:13	475				TART TO SOMETHINGS FAMILS SOME USAGE			8179	***		59.77055	-20.14246
PG354	ļ								L			59.75976	-20.14724
PG355	02/07/98 06:40								S180			59.70844	-20.14479
PG356	02/07/98 06:41	482							\$181			59.70601	-20.14440
PG357	02/07/98 07:00	483							S182			59,66025	-20.13617
PG358	02/07/98 08:00		25						S183			59.51640	-20.11407
PG359	02/07/98 08:52		26						\$184	61		59.42985	-20.13321
PG360	03/07/98 07:08		~~~~~	1	1					62		59.63589	-21.01688
PG361	04/07/98 18:58							T .		63		59.33984	-21.93346
PG362	04/07/98 19:57	1							s	64		59.21224	-21.73355
PG363	04/07/98 21:00	2			1				S	4		59.05168	-21.48510
PG364		1					1			147			
PG365	05/07/98 10:45				1				S185		1	57.13943	-18.63741
<u> </u>			<del></del>		1							<u> </u>	

Sample	Date/ Time (GMT)	chl/pigs	nuts	hydro	halo	N20/CH4	bac	phyto	DMS	sal	Taken by?	lat	lon
PG366	05/07/98 11:38					,,			\$186			57.00775	-18.45381
PG367	05/07/98 12:37								S187			56,86152	-18.24894
PG368	05/07/98 13:41								S188			56.70455	-18.02467
PG369	05/07/98 14:42								S189			56.55622	-17.80527
PG370	05/07/98 15:36								S190	***************************************		56,42102	-17.62000
PG371	05/07/98 16:35								S191			56.27708	-17.41571
PG372	05/07/98 17:33			<u> </u>					S192			56.12899	-17.20925
PG373	05/07/98 18:40								S193			55.95595	-16.97316
PG374	05/07/98 19:57								S194			55.82492	-16,78990
PG375	05/07/98 21:10								\$195			55.64604	-16.52616
PG376	06/07/98 10:53	1							S196			54.11125	-14.45480
PG377	06/07/98 10:57	630		\$1.34								54.10063	-14.44552
PG378	06/07/98 13:14	631		\$2.65					S197			53.78046	-14.00784
PG379	06/07/98 15:04	632		\$3.37					S198			53.52076	-13.66339
PG380	06/07/98 20:16	<b> </b>					ļ			13		52.80463	-12.74414

Polly Machin, BODC

oid	start time	end time	gearcode	comments
AS3058	07/02/98 17:30			Halocarbons
AS3059	07/02/98 18:20			Halocarbons
AS3060	07/02/98 18:40			Halocarbons
AS3061	07/02/98 19:30			Halocarbons
AS3062	07/03/98 02:35			Halocarbons
AS3063	07/03/98 10:34	7777		Halocarbons
AS3064	07/03/98 13:10			Halocarbons
A\$3065	07/03/98 14:14			Halocarbons
AS3066	07/03/98 16:37			Halocarbons
AS3067	07/03/98 20:30			Halocarbons
A\$3068	07/04/98 05:13			Halocarbons
AS3069	07/04/98 09:42			Halocarbons
AS3070	07/04/98 10:11			Halocarbons
A\$3071	07/04/98 13:19			Halocarbons
AS3072	07/04/98 18:37			Halocarbons
AS3073	07/05/98 19:09			Halocarbons
RS1	06/06/98 20:30	07/06/98 05:30	D1	
AS3074	07/06/98 13:47			Halocarbons
A\$3075	07/06/98 19:06			Halocarbons
PG001	09/06/98 09:31			
PG002	09/06/98 09:37			
PG003	09/06/98 10:11			
PG004	09/06/98 10:14			
PG005	09/06/98 11:12			
PG006	09/06/98 12:56			
CTD1	09/06/98 13:47	09/06/98 14:13	13554	
PG007	09/06/98 14:03	00/00/00 1 1.10		
CTD2	09/06/98 15:18	09/06/98 15:40	13555	
CTD3	09/06/98 16:29	09/06/98 17:05	13556	
	10/06/98 06:01	10/06/98 06:38	13557	
CTD4	10/06/98 06:35	10/00/30 00:00	10001	
PG008			<u> </u>	
PG009	10/06/98 06:37			hydrogen peroxide,methylhydroperoxide
AS5001	10/06/98 09:00			Tryatogett potoxiaojinoaryny
PG010	10/06/98 09:00		1	hydrogen peroxide,methylhydroperoxide
AS5002	10/06/98 09:45			hydrogen peroxide,methylhydroperoxide
A\$5003	10/06/98 10:35			nydrogen peronder, med ym y a special
PG011	10/06/98 11:00			
PG012	10/06/98 11:13			
PG013	10/06/98 11:18			hydrogen peroxide,methylhydroperoxide
AS5004	10/06/98 13:00		-	nydrogen peroxide;mearymydroperexide
PG014	10/06/98 13:00			
PG015	10/06/98 13:05			hydrogen peroxide,methylhydroperoxide
AS5005	10/06/98 14:00		1	Tryatogers peroxide, memyimyaroperoxide
PG016	10/06/98 15:00		<del> </del>	
PG017	10/06/98 15:01		1	
PG019	10/06/98 15:03			hydrogen peroxide,methylhydroperoxide
AS5006	10/06/98 16:00		-	hydrogen peroxide,methylhydroperoxide
AS5007	10/06/98 17:00			nyurogen peroxide,mediyinyuroperoxide
PG020	10/06/98 17:03		-	hudragan paravida mathulhudranaravida
AS5008	10/06/98 19:00			hydrogen peroxide,methylhydroperoxide
PG021	10/06/98 19:00			
PG022	10/06/98 21:00			
PG023	10/06/98 22:59			
PG024	11/06/98 01:00			
PG025	11/06/98 03:00			
PG026	11/06/98 05:00		1	
PG027	11/06/98 07:00			
PG028	11/06/98 08:59			
PG029	11/06/98 09:06			
PG030	11/06/98 09:09			
AS5009	11/06/98 10:00			hydrogen peroxide,methylhydroperoxide
PG031	11/06/98 10:58			
AS5010	11/06/98 11:00			hydrogen peroxide,methylhydroperoxide
AS5011	11/06/98 11:30	T		hydrogen peroxide,methylhydroperoxide

oid	start time	end time	gearcode	comments
AS5012	11/06/98 12:35			hydrogen peroxide,methylhydroperoxide
PG032	11/06/98 12:51			
PG033	11/06/98 12:59			
PG034	11/06/98 14:58			
AS5013	11/06/98 15:00			hydrogen peroxide,methylhydroperoxide
AS5014	11/06/98 16:00			hydrogen peroxide,methylhydroperoxide
AS5015	11/06/98 17:00			hydrogen peroxide,methylhydroperoxide
PG035	11/06/98 17:03		<del> </del>	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
PG036	11/06/98 17:06			
AS5016	11/06/98 18:00			hydrogen peroxide,methylhydroperoxide
AS5017	11/06/98 19:00			hydrogen peroxide,methylhydroperoxide
				Hydrogen peroxide, metry mydroperoxide
PG037	11/06/98 19:00			,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,
PG038	11/06/98 19:05			
PG039	11/06/98 21:02			
PG040	11/06/98 23:00			
PG041	12/06/98 01:00			
PG042	12/06/98 03:00			
PG043	12/06/98 05:00			
PG044	12/06/98 07:00			
PG045	12/06/98 08:48			
PG046	12/06/98 08:59			
PG047	12/06/98 09:33			
PG048	12/06/98 10:07			
AS5018	12/06/98 10:20			hydrogen peroxide,methylhydroperoxide
PG049	12/06/98 10:38			
	12/06/98 11:00			hydrogen peroxide,methylhydroperoxide
AS5019			<b></b>	Hydrogen peroxide, meany mydroperoxide
PG050	12/06/98 11:00			
PG051	12/06/98 11:30			L
AS5020	12/06/98 12:00	,.,		hydrogen peroxide,methylhydroperoxide
PG052	12/06/98 12:31			
AS5021	12/06/98 13:00			hydrogen peroxide,methylhydroperoxide
PG053	12/06/98 13:00			
PG055	12/06/98 13:30			
AS5022	12/06/98 14:00			hydrogen peroxide,methylhydroperoxide
PG056	12/06/98 14:32			
AS5023	12/06/98 15:00			hydrogen peroxide,methylhydroperoxide
PG057	12/06/98 15:00			
PG058	12/06/98 15:08			
PG059	12/06/98 16:57			
AS5024	12/06/98 17:00			hydrogen peroxide,methylhydroperoxide
PG060	12/06/98 17:40			Trydrogen peroxide; metry my droperoxide
	.,			
PG061	12/06/98 18:30			
AS1	12/06/98 18:45	12/06/98 19:15		Hydrocarbons
PG062	12/06/98 18:57			
AS5025	12/06/98 19:00			hydrogen peroxide,methylhydroperoxide
PG063	12/06/98 19:03			
PG064	12/06/98 19:40			
AS5026	12/06/98 20:00			hydrogen peroxide,methylhydroperoxide
PG065	12/06/98 20:04			
AS5027	12/06/98 21:00			hydrogen peroxide,methylhydroperoxid
PG066	12/06/98 21:01	ļ		
PG067	12/06/98 23:00			
PG068	13/06/98 01:00			
	<u> </u>			
PG069	13/06/98 03:00			
PG070	13/06/98 05:05		1	
PG071	13/06/98 05:54			
CTD5	13/06/98 07:04	13/06/98 07:55	13558	
PG072	13/06/98 07:15			
PG073	13/06/98 09:40			
PG074	13/06/98 10:05			
PG075	13/06/98 11:32			
CTD6	13/06/98 13:05	13/06/98 14:00	13559	
AS5028	13/06/98 16:00			hydrogen peroxide,methylhydroperoxid
AS5029	13/06/98 17:00	<u> </u>		hydrogen peroxide,methylhydroperoxid

oid	start time	end time	gearcode	comments
CTD7	14/06/98 02:20	14/06/98 02:43	13560	
CTD8	14/06/98 03:15	14/06/98 03:28	13561	
AS5030	14/06/98 04:00			hydrogen peroxide,methylhydroperoxide
CTD9	14/06/98 04:19	14/06/98 04:26	13562	
EAHV1	14/06/98 04:30	14/06/98 15:17		
A\$5031	14/06/98 05:00			hydrogen peroxide,methylhydroperoxide
AS5032	14/06/98 06:00			hydrogen peroxide,methylhydroperoxide
AS5033	14/06/98 07:00			hydrogen peroxide,methylhydroperoxide
AS5034	14/06/98 08:00			hydrogen peroxide,methylhydroperoxide
AS5035	14/06/98 09:00			hydrogen peroxide,methylhydroperoxide
A\$5036	14/06/98 10:00			hydrogen peroxide,methylhydroperoxide
AS5037	14/06/98 11:00			hydrogen peroxide,methylhydroperoxide
AS5038	14/06/98 12:00			hydrogen peroxide,methylhydroperoxide
AS5039	14/06/98 13:00			hydrogen peroxide,methylhydroperoxide
AS5040	14/06/98 14:00			hydrogen peroxide,methylhydroperoxide
AS2	14/06/98 14:28	14/06/98 14:58		Hydrocarbons
AS5041	14/06/98 15:00			hydrogen peroxide,methylhydroperoxide
XBT1	14/06/98 15:12			
PG077	14/06/98 16:03		<u>. ,</u>	
PG078	14/06/98 16:37			
PG079	14/06/98 17:03			
PG079 PG080	14/06/98 17:35			
PG081	14/06/98 18:05			
PG082	14/06/98 18:35			
PG083	14/06/98 19:02			
PG084	14/06/98 19:32			
PG085	14/06/98 20:07			
	14/06/98 20:38			
PG086	14/06/98 21:00			
PG087				
PG088	14/06/98 21:30			
PG089	14/06/98 22:01			
PG090	14/06/98 22:31			
PG091	14/06/98 23:00			
PG092	14/06/98 23:30			
PG093	14/06/98 23:59			
PG094	15/06/98 00:30			
PG095	15/06/98 01:00			
PG096	15/06/98 01:25			
PG097	15/06/98 02:23		40500	
CTD10	15/06/98 02:27	15/06/98 02:51	13563	
CTD11	15/06/98 03:23	15/06/98 03:43	13564	
AS5042	15/06/98 04:00			hydrogen peroxide,methylhydroperoxide
CTD12	15/06/98 04:08	15/06/98 04:15	13565	
EAHV2	15/06/98 04:25	15/06/98 14:48		t. D. d
AS5043	15/06/98 05:00			hydrogen peroxide, methylhydroperoxide
A\$5044	15/06/98 06:00			hydrogen peroxide,methylhydroperoxide
AS5045	15/06/98 07:00			hydrogen peroxide,methylhydroperoxide
AS5046	15/06/98 08:00			hydrogen peroxide,methylhydroperoxide
AS5047	15/06/98 09:00			hydrogen peroxide,methylhydroperoxide
A\$5048	15/06/98 10:00			hydrogen peroxide,methylhydroperoxide
AS5049	15/06/98 11:00			hydrogen peroxide,methylhydroperoxide
A\$5050	15/06/98 12:00			hydrogen peroxide,methylhydroperoxide
AS5051	15/06/98 13:00			hydrogen peroxide,methylhydroperoxide
AS5052	15/06/98 14:00			hydrogen peroxide,methylhydroperoxide
AS3	15/06/98 14:16	15/06/98 15:02		Hydrocarbons
AS5053	15/06/98 15:00			hydrogen peroxide,methylhydroperoxide
CTD13	15/06/98 15:03	15/06/98 15:30	13566	
PG098	15/06/98 16:00			
PG099	15/06/98 17:05			
PG100	15/06/98 18:01			
PG101	15/06/98 19:02			
PG102	15/06/98 20:06		-	
PG103	15/06/98 21:01			
	15/06/98 21:33	<del> </del>		

oid	start time	end time	gearcode	comments
PG105	15/06/98 22:02			
PG106	15/06/98 23:00			
PG107	15/06/98 23:30			
PG108	16/06/98 00:01			
PG109	16/06/98 00:30			
PG110	16/06/98 01:01			
PG111	16/06/98 01:30			
PG112	16/06/98 02:00			
CTD14	16/06/98 02:15	16/06/98 02:35	13567	
CTD15	16/06/98 03:05	16/06/98 03:20	13568	
CTD16	16/06/98 04:01	16/06/98 04:09	13569	
EAHV3	16/06/98 04:33	16/06/98 13:40		
AS5054	16/06/98 06:15			hydrogen peroxide,methylhydroperoxide
AS5055	16/06/98 07:00			hydrogen peroxide,methylhydroperoxide
AS5056	16/06/98 08:00			hydrogen peroxide,methylhydroperoxide
A\$5057	16/06/98 09:00			hydrogen peroxide,methylhydroperoxide
AS5058	16/06/98 10:00			hydrogen peroxide,methylhydroperoxide
A\$5059	16/06/98 11:00		****	hydrogen peroxide,methylhydroperoxide
AS5060	16/06/98 12:00	†		hydrogen peroxide,methylhydroperoxide
AS5061	16/06/98 13:00			hydrogen peroxide,methylhydroperoxide
AS5062	16/06/98 14:00			hydrogen peroxide,methylhydroperoxide
AS5063	16/06/98 15:00			
PG113	16/06/98 16:00	<del>                                     </del>		hydrogen peroxide,methylhydroperoxide
PG114	16/06/98 16:30			
PG115	16/06/98 17:00		77 748 444 444 444 444 444 444 444 444 4	
PG116	16/06/98 18:00			
PG117	16/06/98 18:35			
PG118	16/06/98 19:00	ļ		
PG119	16/06/98 19:30	<u> </u>		
PG120	16/06/98 20:00			
PG121		-		
PG121	16/06/98 20:30 16/06/98 21:00			, , , , , , , , , , , , , , , , , , ,
PG123	16/06/98 21:30			
PG124	16/06/98 22:00			
PG125	16/06/98 22:31			
PG126	16/06/98 23:03			
PG127	16/06/98 23:30			
PG128	17/06/98 00:01			
PG139	17/06/98 00:01			
PG129	17/06/98 00:30			
PG130	17/06/98 01:01			
PG131	17/06/98 01:35			
PG132	17/06/98 01:59			
CTD17	17/06/98 02:14	17/06/98 02:35	13570	
CTD18	17/06/98 03:09	17/06/98 03:24	13571	
CTD19	17/06/98 04:15	17/06/98 04:22	13572	
EAHV4	17/06/98 04:30	17/06/98 20:43		
AS5064	17/06/98 05:00			hydrogen peroxide,methylhydroperoxide
AS5065	17/06/98 06:00			hydrogen peroxide, methylhydroperoxide
AS5066	17/06/98 07:00			hydrogen peroxide,methylhydroperoxide
AS5067	17/06/98 08:00			hydrogen peroxide,methylhydroperoxide
AS5068	17/06/98 09:00			hydrogen peroxide,methylhydroperoxide
AS5069	17/06/98 10:00			hydrogen peroxide,methylhydroperoxide
AS5070	17/06/98 11:00			hydrogen peroxide,methylhydroperoxide
AS5071	17/06/98 12:00		7/7-201/	hydrogen peroxide,methylhydroperoxide
AS5072	17/06/98 13:00			hydrogen peroxide,methylhydroperoxide
AS5073	17/06/98 14:00			hydrogen peroxide,methylhydroperoxide
AS4	17/06/98 14:26	17/06/98 15:31		Hydrocarbons
AS5074	17/06/98 15:00			hydrogen peroxide,methylhydroperoxide
AS5075	17/06/98 16:00		· · · · · · · · · · · · · · · · · · ·	hydrogen peroxide,methylhydroperoxide
BOT1	17/06/98 16:15	17/06/98 16:20	13573	ar agon ker ayide imeni hin har ober ayide
				harden and many file and the file of the file
AS5076	17/06/98 17:00 1	1		
AS5076 AS5077	17/06/98 17:00 17/06/98 18:00			hydrogen peroxide,methylhydroperoxide hydrogen peroxide,methylhydroperoxide

oid	start time	end time	gearcode	comments
AS5079	17/06/98 20:00			hydrogen peroxide,methylhydroperoxide
AS5080	17/06/98 21:00			hydrogen peroxide,methylhydroperoxide
PG133	17/06/98 21:10			
PG134	17/06/98 21:30			hydrogon porovido mothylhydroperovido
AS5081	17/06/98 22:00			hydrogen peroxide,methylhydroperoxide
PG135	17/06/98 22:00			
PG136	17/06/98 22:30	100000000	D0	
RS2	17/06/98 22:30	18/06/98 06:15	D2	
PG137	17/06/98 23:00			
PG138	17/06/98 23:30			
PG140	18/06/98 00:30			
PG141	18/06/98 01:00			
PG142	18/06/98 01:30			
PG143	18/06/98 01:34			
PG144	18/06/98 02:00			
PG145	18/06/98 02:30		10571	
CTD20	18/06/98 03:03	18/06/98 03:22	13574	
BOT2	18/06/98 03:53	18/06/98 03:55	13575	
CTD21	18/06/98 04:11	18/06/98 04:24	13576	
EAHV5	18/06/98 04:30	18/06/98 15:15		to de la constitución de la cons
AS5082	18/06/98 07:00			hydrogen peroxide,methylhydroperoxide
AS5083	18/06/98 08:00			hydrogen peroxide, methylhydroperoxide
AS5084	18/06/98 09:00			hydrogen peroxide,methylhydroperoxide
AS5085	18/06/98 10:00			hydrogen peroxide,methylhydroperoxide
AS5086	18/06/98 11:00			hydrogen peroxide, methylhydroperoxide
A\$5087	18/06/98 12:00			hydrogen peroxide,methylhydroperoxide
AS5088	18/06/98 13:00			hydrogen peroxide,methylhydroperoxide
AS5089	18/06/98 14:00		··-	hydrogen peroxide,methylhydroperoxide
AS1001	18/06/98 14:10			DMS
A\$5090	18/06/98 15:00			hydrogen peroxide,methylhydroperoxide
AS1002	18/06/98 15:32			DMS
AS5091	18/06/98 16:00			hydrogen peroxide, methylhydroperoxide
PG146	18/06/98 16:03			
PG147	18/06/98 16:31			
PG148	18/06/98 17:00			
PG149	18/06/98 17:30			
PG150	18/06/98 18:34			
PG151	18/06/98 19:00			
PG152	18/06/98 19:20			
PG153	18/06/98 19:32			
PG154	18/06/98 20:02			
PG155	18/06/98 20:31			
PG156	18/06/98 21:01			
PG157	18/06/98 21:31			
PG158	18/06/98 21:43			
PG159	18/06/98 22:33			
PG159	18/06/98 23:00			
PG160	18/06/98 23:31			
PG162	19/06/98 00:00			
PG162	19/06/98 00:31			
PG164	19/06/98 01:00			
PG165	19/06/98 01:10			
PG166	19/06/98 01:55			
CTD22	19/06/98 02:39	19/06/98 02:46	13577	
R\$3	19/06/98 06:30	19/06/98 10:30	D3	
AS5092	19/06/98 09:00	15/50/50 10.50	50	hydrogen peroxide,methylhydroperoxide
	19/06/98 10:00		<del> </del>	hydrogen peroxide,methylhydroperoxide
A\$5093	19/06/98 10:00			illana San Barayina ilina ilini arabarayina
PG167	19/06/98 10:30			hydrogen peroxide,methylhydroperoxide
AS5094		<b> </b>		hydrogen peroxide,methylhydroperoxide
AS5095	19/06/98 13:00	10/06/09 21:20	D4	ilyarogen peroniae,memyinyaroperoniae
RS4	19/06/98 15:00	19/06/98 21:30	U4	hydrogen peroxide,methylhydroperoxide
A\$5096	19/06/98 19:00	20/00/00 43:00	D5	Hydrogen peroxide, methylinguroperoxide
RS5	20/06/98 09:00	20/06/98 13:00	UO	hydrogen peroxide,methylhydroperoxide

oid	start time	end time	gearcode	comments
AS5098	20/06/98 12:00			hydrogen peroxide,methylhydroperoxide
AS5099	20/06/98 13:00	.,	· · · · · · · · · · · · · · · · · · ·	hydrogen peroxide,methylhydroperoxide
AS5100	20/06/98 14:00			hydrogen peroxide,methylhydroperoxide
PG168	20/06/98 15:08			
PG169	20/06/98 15:30			
AS5101	20/06/98 16:00			hydrogen peroxide,methylhydroperoxide
PG170	20/06/98 16:03			
AS1003	20/06/98 16:15			DMS
AS5	20/06/98 16:57	20/06/98 17:40	A Province Control Con	Hydrocarbons
PG171	20/06/98 17:00		······································	
PG172	20/06/98 18:00			<u> </u>
PG173	20/06/98 19:00			,
PG174	20/06/98 19:32			
PG175	20/06/98 20:00			
PG176	20/06/98 20:03		V	
PG177	20/06/98 20:30			
PG178	20/06/98 21:00			
CTD23	20/06/98 22:09	20/06/98 22:33	13578	
PG179	20/06/98 22:28	20/00/00 22:00	,,,,,,	
AS6	21/06/98 00:46	21/06/98 00:59		Hydrocarbons
AS7	21/06/98 01:56	21/06/98 02:31	· · · · · · · · · · · · · · · · · · ·	Hydrocarbons
CTD24	21/06/98 03:02	21/06/98 03:20	13579	Trydrocarpons
CTD24	21/06/98 03:47	21/06/98 03:59	13580	1
	21/06/98 04:00	21/00/90 00:39	13300	hydrogen peroxide,methylhydroperoxide
AS5102				<u> </u>
AS1004	21/06/98 04:04			DMS
EAHV6	21/06/98 04:27	21/06/98 14:47		Ma 12
A\$1005	21/06/98 05:00		<del></del>	DMS
AS5103	21/06/98 05:00			hydrogen peroxide,methylhydroperoxide
AS1006	21/06/98 06:00			DMS
AS5104	21/06/98 06:00			hydrogen peroxide,methylhydroperoxide
AS1007	21/06/98 07:00			DMS
AS5105	21/06/98 07:00			hydrogen peroxide,methylhydroperoxide
AS1008	21/06/98 08:00			DMS
AS5106	21/06/98 08:00			hydrogen peroxide,methylhydroperoxide
AS1009	21/06/98 09:00			DMS
AS5107	21/06/98 09:00			hydrogen peroxide,methylhydroperoxide
AS1010	21/06/98 10:00			DMS
AS5108	21/06/98 10:00			hydrogen peroxide,methylhydroperoxide
AS1011	21/06/98 10:59			DMS
AS5109	21/06/98 11:00			hydrogen peroxide,methylhydroperoxide
AS1012	21/06/98 11:59			DMS
AS5110	21/06/98 12:00			hydrogen peroxide,methylhydroperoxide
AS1013	21/06/98 12:59			DMS
AS5111	21/06/98 13:00			hydrogen peroxide,methylhydroperoxide
AS1014	21/06/98 14:00			DMS
AS5112	21/06/98 14:00			hydrogen peroxide,methylhydroperoxide
AS8	21/06/98 14:56	21/06/98 15:51		Hydrocarbons
AS1015	21/06/98 15:00			DMS
AS5113	21/06/98 15:00		*	hydrogen peroxide,methylhydroperoxide
CTD26	21/06/98 15:05	21/06/98 15:33	13581	An again has a season and an about of the
AS1016	21/06/98 15:50		, +041	DMS
AS5114	21/06/98 16:00	-		hydrogen peroxide,methylhydroperoxide
BOT3	21/06/98 16:15	21/06/98 16:20	13582	nyaragan peroxido, menyinyaraperoxide
PG180	21/06/98 19:00	21100100 10.20	1002	
PG181	21/06/98 19:33			
PG182	21/06/98 20:00	-		
PG183	21/06/98 20:30			
PG184	21/06/98 21:00			
PG185	21/06/98 21:30			
EAHV7	21/06/98 21:42	22/06/98 08:53		
AS9	22/06/98 00:02	22/06/98 00:48		Hydrocarbons
AS1017	22/06/98 09:00		· · · · · · · · · · · · · · · · · · ·	DMS
AS1018	22/06/98 09:20			DMS

old	start time	end time	gearcode	comments
AS5115	22/06/98 09:20			hydrogen peroxide,methylhydroperoxide
AS1019	22/06/98 09:58			DMS
AS5116	22/06/98 10:00			hydrogen peroxide,methylhydroperoxide
AS1020	22/06/98 10:53			DMS
AS5117	22/06/98 11:00			hydrogen peroxide,methylhydroperoxide
CTD27	22/06/98 12:00	22/06/98 13:00	13583	
AS5118	22/06/98 12:00			hydrogen peroxide,methylhydroperoxide
AS5119	22/06/98 13:00			hydrogen peroxide,methylhydroperoxide
BOT4	22/06/98 13:05	22/06/98 13:10	13584	
		2200,00 10.10	7000 1	DMS
AS1021	22/06/98 13:37 22/06/98 14:00			hydrogen peroxide,methylhydroperoxide
AS5120				Trydrogen peroxide, memyling aroperoxide
PG186	22/06/98 14:15			DMS
AS1022	22/06/98 14:22			hydrogen peroxide,methylhydroperoxide
AS5121	22/06/98 15:00			nydrogen peroxide,metriyiriydroperoxide
PG187	22/06/98 15:15			
XBT2	22/06/98 15:30			
PG188	22/06/98 16:00			
PG189	22/06/98 17:00		A-04 V 3 V V V V V V V	
PG190	22/06/98 18:15			
PG191	22/06/98 18:44			
PG192	22/06/98 19:00		·····	
PG193	22/06/98 19:36			
PG194	22/06/98 20:01			
PG195	22/06/98 20:35			
PG196	22/06/98 21:02			
PG197	22/06/98 21:30			
PG198	22/06/98 22:00			
PG199	22/06/98 22:30			
PG200	22/06/98 23:02			
PG201	22/06/98 23:30			
PG202	23/06/98 00:01			
PG203	23/06/98 00:31			
PG204	23/06/98 01:00			
PG205	23/06/98 01:31			
PG206	23/06/98 02:00			
PG207	23/06/98 02:32			
CTD28	23/06/98 02:53	23/06/98 02:59	13585	
CTD29	23/06/98 03:22	23/06/98 03:42	13586	
PG208	23/06/98 03:35			
AS5122	23/06/98 04:00			hydrogen peroxide,methylhydroperoxide
AS5123	23/06/98 05:00			hydrogen peroxide,methylhydroperoxide
CTD30	23/06/98 05:58	23/06/98 06:17	13587	
AS5124	23/06/98 06:00			hydrogen peroxide,methylhydroperoxide
AS5125	23/06/98 07:00			hydrogen peroxide,methylhydroperoxide
RS6	23/06/98 07:30	23/06/98 15:00	D7 and D8	
CTD31	23/06/98 08:05	23/06/98 08:20	13588	
A\$5126	23/06/98 10:00			hydrogen peroxide,methylhydroperoxide
CTD32	23/06/98 10:23	23/06/98 10:43	13589	
AS5127	23/06/98 11:00			hydrogen peroxide,methylhydroperoxide
AS5128	23/06/98 12:00			hydrogen peroxide,methylhydroperoxide
вот5	23/06/98 13:04	23/06/98 13:06	13589	
CTD33	23/06/98 13:11	23/06/98 13:35	13591	
вот6	24/06/98 16:50	24/06/98 16:55	13592	
RS7	24/06/98 21:30	24/06/98 23:15	D10	
PG209	25/06/98 00:01			
PG210	25/06/98 00:21			
RS8	25/06/98 00:30	25/06/98 09:00	D11	
PG211	25/06/98 00:41			
PG212	25/06/98 01:00			
PG213	25/06/98 01:20			
PG214	25/06/98 01:39			
PG214	25/06/98 02:00	<del>                                     </del>		
PG216	25/06/98 02:16			

oid	start time	end time	gearcode	comments
PG218	25/06/98 02:45			
PG219	25/06/98 03:00			
вот7	25/06/98 11:15	25/06/98 11:30	13593	
BK1	25/06/98 11:20		SEB7	DMSP lyase
AS5129	25/06/98 12:00			hydrogen peroxide,methylhydroperoxide
AS5130	25/06/98 13:00			hydrogen peroxide,methylhydroperoxide
AS1023	25/06/98 13:40			DMS
AS5131	25/06/98 14:00			hydrogen peroxide,methylhydroperoxide
PG220	25/06/98 14:00			Tryarogett peroxide, metryinyaroperoxide
EAHV8	25/06/98 14:30		26/06/98 03:40	
AS1024	25/06/98 14:53		201000000000000000000000000000000000000	DMC
AS5132	25/06/98 15:00			DMS
PG221	25/06/98 15:04			hydrogen peroxide,methylhydroperoxide
PG222	25/06/98 15:35	<u></u>		
BK2	25/06/98 16:00		SEB8	DMCD hase
BOT8	25/06/98 16:03	25/06/98 16:05	13594	DMSP lyase
PG223	25/06/98 16:34	23/00/90 10:05	13094	
AS1025	25/06/98 16:35		ļ	DMS
AS5133	25/06/98 17:00	<del> </del>		hydrogen peroxide,methylhydroperoxide
PG224	25/06/98 17:00			
AS1026	25/06/98 17:07			DMS
AS10	25/06/98 17:29	25/06/98 18:10		Hydrocarbons
AS1027	25/06/98 17:37			DMS
AS5134	25/06/98 18:00			hydrogen peroxide, methylhydroperoxide
PG225	25/06/98 18:00			
AS1028	25/06/98 18:04			DMS
PG226	25/06/98 18:30			Diffo
AS5135	25/06/98 19:00			hydrogen peroxide,methylhydroperoxide
вот9	25/06/98 19:01	25/06/98 19:03	13595	nydrogen peroxide, metrry mydroperoxide
BK3	25/06/98 19:02	20/00/30 19:00	SEB9	DMSP lyase
PG227	25/06/98 19:31		OLDS	DIVIOR IYase
AS1029		<b></b>		DNIO
	25/06/98 19:40			DMS
AS5136	25/06/98 20:00	<u> </u>		hydrogen peroxide,methylhydroperoxide
PG228	25/06/98 20:01			
PG229	25/06/98 20:04			
AS1030	25/06/98 20:09			DMS
PG230	25/06/98 20:58		- 3 - 5 - 5 - 5 - 5 - 5 - 5 - 5 - 5 - 5	
AS5137	25/06/98 21:00			hydrogen peroxide, methylhydroperoxide
AS1031	25/06/98 21:00			DMS
PG231	25/06/98 21:32			
AS5138	25/06/98 22:00			hydrogen peroxide,methylhydroperoxide
BOT10	25/06/98 22:00	25/06/98 22:02	13596	
BK4	25/06/98 22:02		SEB10	DMSP lyase
AS1032	25/06/98 22:15			DMS
PG232	25/06/98 22:32			
AS5139	25/06/98 23:00			hydrogen peroxide,methylhydroperoxide
AS1033	25/06/98 23:00			DMS
PG233	25/06/98 23:00		<del> </del>	DINO
PG234	25/06/98 23:30		<u> </u>	
			-	harden and harden state of the
AS5140	26/06/98 00:00			hydrogen peroxide,methylhydroperoxide
AS1034	26/06/98 00:00			DMS
PG235	26/06/98 00:01			· · · · · · · · · · · · · · · · · · ·
AS5141	26/06/98 01:00			hydrogen peroxide,methylhydroperoxide
AS1035	26/06/98 01:02			DMS
AS5142	26/06/98 02:00			hydrogen peroxide,methylhydroperoxide
AS1036	26/06/98 02:00			DMS
EAHV9	26/06/98 10:30	26/06/98 18:23		
ХВТЗ	26/06/98 13:03			
XBT4	26/06/98 14:01			
XBT5	26/06/98 14:57		<del>                                     </del>	
	26/06/98 16:05		<del> </del>	
XBT6	Z0/U0/90 10.UD	1	1 ,	

oid	start time	end time	gearcode	comments
XBT8	26/06/98 18:04			
хвт9	26/06/98 19:01			
PG236	26/06/98 19:55			
XBT10	26/06/98 19:57			
XBT11	26/06/98 20:55			
CTD34	26/06/98 21:05	26/06/98 21:30	13598	
BOT11	26/06/98 21:06	26/06/98 21:08	13597	
PG237	26/06/98 21:25			
PG238	27/06/98 00:02			
CTD35	27/06/98 02:18	27/06/98 02:41	13599	
BOT12	27/06/98 03:08	27/06/98 03:10	13600	
	27/06/98 03:25	27/06/98 03:45	13601	
CTD36	27/06/98 03:59	21700/00 00: 10		
PG239	27/06/98 19:14	27/06/98 20:05		Hydrocarbons
AS11	27/06/98 21:30	27/06/98 21:45	13602	
CTD37		27/06/98 21:55	13603	
BOT13	27/06/98 21:50	2/100/90 21.50	10000	
PG240	27/06/98 23:02			
PG241	28/06/98 00:08			
PG242	28/06/98 01:30	26108106 V3-33	13604	
CTD38	28/06/98 02:09	28/06/98 02:33	13605	
CTD39	28/06/98 03:18	28/06/98 03:41	13000	
PG243	28/06/98 04:08	00/00/00 45:00	12606	
CTD40	28/06/98 14:19	28/06/98 15:02	13606	
PG244	28/06/98 16:00			40.40 104.45
EAHV10	28/06/98 16:06	29/06/98 08:57		not between 16:40 and 21:45
AS5143	28/06/98 16:10			hydrogen peroxide,methylhydroperoxide
PG245	28/06/98 17:04			
PG246	28/06/98 18:04			
PG247	28/06/98 19:00			
PG248	28/06/98 20:00			
PG249	28/06/98 20:59			
AS5144	28/06/98 21:00			hydrogen peroxide,methylhydroperoxide
AS5145	28/06/98 22:00			hydrogen peroxide, methylhydroperoxide
PG250	28/06/98 22:00			
AS5146	28/06/98 23:00			hydrogen peroxide,methylhydroperoxide
PG251	28/06/98 23:00			
AS5147	29/06/98 00:00			hydrogen peroxide,methylhydroperoxide
PG252	29/06/98 00:00			
AS12	29/06/98 00:45			Hydrocarbons
AS5148	29/06/98 01:00			hydrogen peroxide,methylhydroperoxide
PG253	29/06/98 01:00			
A\$13	29/06/98 01:05	29/06/98 01:47		Hydrocarbons
AS5149	29/06/98 02:00			hydrogen peroxide,methylhydroperoxide
PG254	29/06/98 02:00			hydrogen peroxide,methylhydroperoxide
AS5150	29/06/98 03:00			Hydrogen peroxide, meally mydroperoxide
PG255	29/06/98 03:00		-	
AS5151	29/06/98 04:00			hydrogen peroxide,methylhydroperoxide
PG256	29/06/98 04:00			
AS5152	29/06/98 05:00			hydrogen peroxide,methylhydroperoxide
PG257	29/06/98 05:00	1	1	
PG258	29/06/98 06:00	-	<del></del>	
PG259	29/06/98 07:00			
PG260	29/06/98 08:03			
PG261	29/06/98 09:00			
EAHV11	29/06/98 09:31	29/06/98 16:52		
PG262	29/06/98 10:00			
PG263	29/06/98 11:00			
PG264				
PG265	29/06/98 13:00			
AS5153			1	hydrogen peroxide,methylhydroperoxide
	29/06/98 14:00			To the state of th
PG266	/M/IN/MA IZLIII			

oid	start time	end time	gearcode	comments
PG267	29/06/98 15:00			
PG268	29/06/98 15:03			
PG269	29/06/98 15:59			
AS5155	29/06/98 16:00			hydrogen peroxide,methylhydroperoxide
PG270	29/06/98 16:02			
PG271	29/06/98 16:58			
AS5156	29/06/98 17:00			hydrogen peroxide,methylhydroperoxide
PG272	29/06/98 17:00			
AS14	29/06/98 17:33	29/06/98 18:13		Hydrocarbone
		28/00/90 10.13		Hydrocarbons
AS5157	29/06/98 18:00			hydrogen peroxide,methylhydroperoxide
PG273	29/06/98 18:00			
PG274	29/06/98 18:35			
PG275	29/06/98 18:38			
AS15	29/06/98 18:54			Hydrocarbons
AS5158	29/06/98 19:00			hydrogen peroxide,methylhydroperoxide
PG276	29/06/98 19:01	<u> </u>		Tryalogen peroxide; meany my droper oxide
ļ ———		00/00/00 (0.05		
EAHV12	29/06/98 19:58	30/06/98 10:35	ļ	
AS5159	29/06/98 20:00			hydrogen peroxide, methylhydroperoxide
PG277	29/06/98 20:01			
PG279	29/06/98 21:00			
PG278	29/06/98 21:02			
AS5160	29/06/98 22:00			hydrogen peroxide,methylhydroperoxide
PG280	29/06/98 22:00			
PG281	29/06/98 23:00			
PG282	30/06/98 00:00			
PG283	30/06/98 01:00			
PG284	30/06/98 02:00			
PG285	30/06/98 03:00			
PG286	30/06/98 04:00			
PG287	30/06/98 05:00			
PG288	30/06/98 06:00			
PG289	30/06/98 07:00			
PG290	30/06/98 08:00			
PG291	30/06/98 09:00			
PG292	30/06/98 09:47	***************************************		
PG293	30/06/98 10:00			
PG294	30/06/98 11:00			
PG295	30/06/98 12:00			
PG296	30/06/98 13:00	1		
PG297	30/06/98 13:03			
PG298	30/06/98 14:00		-	
PG299	30/06/98 15:00			
PG300	30/06/98 16:00			<u></u>
PG301	30/06/98 16:30			
PG302	30/06/98 17:00			
PG303	30/06/98 17:59			
PG304	30/06/98 18:00		<u> </u>	
PG305	30/06/98 19:00			
AS16	30/06/98 19:35		<del> </del>	Hydrocarbons
PG306	30/06/98 20:00			Trydrodizono
PG307	30/06/98 20:09	1		
		20/00/00 04:00		115
A\$17	30/06/98 20:25	30/06/98 21:03		Hydrocarbons
PG308	30/06/98 21:00		<b> </b>	
EAHV13	30/06/98 22:00	01/07/98 07:00		
PG309	30/06/98 22:00			
PG310	30/06/98 23:00			
PG311	01/07/98 00:00		~~~	· · · · · · · · · · · · · · · · · · ·
PG312		<del>                                     </del>		
	01/07/98 01:00	<u> </u>	ļ	<u> </u>
PG313	01/07/98 02:00			
PG314	01/07/98 03:00			
PG315	01/07/98 04:00			

oid	start time	end time	gearcode	comments
	01/07/98 05:00		<u> </u>	
PG316				
PG317A	01/07/98 06:00			
PG317	01/07/98 09:05			
PG318	01/07/98 10:00			
PG319	01/07/98 11:00			
PG320	01/07/98 11:50			
PG321	01/07/98 13:01			
PG322	01/07/98 14:00			
	01/07/98 14:48	01/07/98 22:38		
EAHV14	01/07/98 15:00	01/01/90 22:50		
PG323	01/07/98 15:06			
PG324	01/07/98 15:42			
PG325	01/07/98 16:00			hydrogen peroxide,methylhydroperoxide
AS5161	01/07/98 16:00			nya.ogon polonari
PG326				
PG327	01/07/98 16:59			DMS
AS1037	01/07/98 17:00			hydrogen peroxide,methylhydroperoxide
AS5162	01/07/98 17:00			riyarogen peroxide,metriyinydroperoxide
PG328	01/07/98 17:00			hydrogen peroxide,methylhydroperoxide
A\$5163	01/07/98 18:00			пуагоден регохіае, нешулкуагорегохіае
PG329	01/07/98 18:04			
PG330	01/07/98 19:00			hydrogon porovido mothylhydronorovido
AS5164	01/07/98 19:00			hydrogen peroxide,methylhydroperoxide
PG331	01/07/98 19:20			L. L
AS5165	01/07/98 20:00			hydrogen peroxide,methylhydroperoxide
PG332	01/07/98 20:02			
PG333	01/07/98 20:59			the state of the s
AS5166	01/07/98 21:00			hydrogen peroxide,methylhydroperoxide
PG334	01/07/98 21:01			
PG335	01/07/98 21:10			
A\$1038	01/07/98 21:40			DMS
PG336	01/07/98 21:46			
PG337	01/07/98 21:50			
PG338	01/07/98 22:00			
PG339	01/07/98 22:02			
PG340	01/07/98 23:00			
PG341	01/07/98 23:17			
PG342	02/07/98 00:00			
PG343	02/07/98 01:00			
PG344	02/07/98 02:00			
PG345	02/07/98 03:00			
PG346	02/07/98 04:00	-		
PG347	02/07/98 05:00			
PG348	02/07/98 05:14			
PG349	02/07/98 05:15	<del> </del>		
PG350	02/07/98 05:41	<u> </u>	1	
PG351	02/07/98 05:42			
PG352	02/07/98 06:00			
PG353	02/07/98 06:13			
PG354	02/07/98 06:18		1	
				DMS
A\$1039				
PG355	02/07/98 06:40		_	
PG356	02/07/98 06:41			
PG357	02/07/98 07:00			
A\$1040	02/07/98 07:24			DMS
PG358	02/07/98 08:00			
AS1041				DMS
PG359	02/07/98 08:52			
AS5167				hydrogen peroxide,methylhydroperoxide
		-		DMS
AS1042				DMS
AS1043				
AS1044	02/07/98 12:07		1	DMS

oid	start time	end time	gearcode	comments
AS1045	02/07/98 12:30			DMS
AS5168	02/07/98 14:00			hydrogen peroxide,methylhydroperoxide
AS1046	02/07/98 14:08			DMS
AS5169	02/07/98 15:00			hydrogen peroxide,methylhydroperoxide
AS5170	02/07/98 16:00			hydrogen peroxide,methylhydroperoxide
AS5171	02/07/98 17:00			hydrogen peroxide,methylhydroperoxide
AS1047	02/07/98 17:04			DMS
CTD41	02/07/98 17:15	02/07/98 17:30	13607	
CTD42	02/07/98 17:40	02/07/98 19:40	13608	
AS1048	02/07/98 17:45			DMS
AS5172	02/07/98 18:00			hydrogen peroxide,methylhydroperoxide
BK5	02/07/98 18:36		CTD42	Microbial diversity
AS1049	02/07/98 18:43			DMS
AS5173	02/07/98 19:00			hydrogen peroxide,methylhydroperoxide
AS1050	02/07/98 19:36		<u></u>	DMS
AS1051	02/07/98 20:40			DMS
AS1052	02/07/98 21:15			DMS
CTD43	02/07/98 21:26	02/07/98 23:37	13609	
AS1053	02/07/98 21:55			DMS
BK6	02/07/98 22:16		CTD43	Microbial diversity
CTD44	03/07/98 01:20	03/07/98 03:06	13610	
BK7	03/07/98 03:18		CTD44	Microbial diversity
CTD45	03/07/98 04:40	03/07/98 06:34	13611	
BK8	03/07/98 06:51		CTD45	Microbial diversity
PG360	03/07/98 07:08		~ FA	
CTD46	03/07/98 08:14	03/07/98 10:14	13612	
BK9	03/07/98 09:32		CTD46	Microbial diversity
CTD47	03/07/98 11:42	03/07/98 13:37	13613	
CTD48	03/07/98 14:53	03/07/98 16:46	13614	
BK10	03/07/98 16:56		CTD48	Microbial diversity
CTD49	03/07/98 18:22	03/07/98 20:15	13615	
BK11	03/07/98 19:59		CTD49	Microbial diversity
CTD50	03/07/98 22:20	04/07/98 00:07	13616	
BK12	03/07/98 23:30		CTD50	Microbial diversity
CTD51	04/07/98 04:42	04/07/98 04:52	13617	
CTD52	04/07/98 05:15	04/07/98 05:34	13618	
EAHV15	04/07/98 05:37	04/07/98 17:51		
RS9	04/07/98 06:00	04/07/98 13:00	D12	
RS10	04/07/98 16:45	04/07/98 18:42	D13	
CTD53	04/07/98 18:34	04/07/98 18:42	13619	
PG361	04/07/98 18:58			
PG362	04/07/98 19:57			
PG363	04/07/98 21:00	05035555	40000	
BOT14	05/07/98 07:04	05/07/98 07:07	13620	<u> </u>
EAHV16	05/07/98 10:28	05/07/98 19:07		
PG365	05/07/98 10:45			
PG366	05/07/98 11:38	Am (Am		
RS11	05/07/98 12:30	05/07/98 13:30	D14	
PG367	05/07/98 12:37			
PG368	05/07/98 13:41			
PG369	05/07/98 14:42			
PG370	05/07/98 15:36			
PG371	05/07/98 16:35			
PG372	05/07/98 17:33	<u> </u>		
PG373	05/07/98 18:40	0503504554	4000	
BOT15	05/07/98 19:02	05/07/98 19:04	13621	
PG374	05/07/98 19:57			
PG375	05/07/98 21:10	06/07/09 00:05	10000	
CTD54	06/07/98 06:12	06/07/98 08:35	13622	
BOT16	06/07/98 08:41	06/07/98 08:42	13623	
PG376 PG377	06/07/98 10:53			
PG377	06/07/98 10:57			<u> </u>
L G210	06/07/98 13:14			

oid	start time	end time	gearcode	comments
PG379	06/07/98 15:04			
BOT17	06/07/98 19:00	06/07/98 19:01	13624	
PG380	06/07/98 20:16			
BOT18	07/07/98 05:57	07/07/98 05:58	13625	
\S3076	07/07/98 07:32			Halocarbons
BOT19	07/07/98 17:57	07/07/98 17:58	13626	
AS3077	07/07/98 19:09			Halocarbons
AS3001	06/10/98 06:40			Halocarbons
A\$3002	06/10/98 16:48			Halocarbons
AS3003	06/10/98 22:20			Halocarbons
A\$3004	06/11/98 08:50			Halocarbons
A\$3005	06/11/98 12:35			Halocarbons
AS3006	06/11/98 14:40			Halocarbons
AS3007	06/12/98 10:44			Halocarbons
AS3008	06/12/98 14:27			Halocarbons
AS3009	06/12/98 19:41			Halocarbons
AS3010	13/06/98 12:48			Halocarbons
AS3011	14/06/98 01:53			Halocarbons
A\$3012	14/06/98 07:10			Halocarbons
	14/06/98 08:10			Haiocarbons
AS3013	14/06/98 08:40			Halocarbons
AS3014	14/06/98 09:10			Halocarbons
AS3015				Halocarbons
AS3016	14/06/98 09:40			Halocarbons
AS3017	14/06/98 14:40			Halocarbons
AS3018	15/06/98 13:13			Halocarbons
AS3019	15/06/98 13:43			Halocarbons
A\$3020	15/06/98 14:14	ļ		Halocarbons
A\$3021	15/06/98 22:35			Halocarbons
AS3022	16/06/98 02:39			Halocarbons
AS3023	16/06/98 13:07			Halocarbons
A\$3024	16/06/98 13:37			Halocarbons
AS3025	16/06/98 22:03			Halocarbons
AS3027	17/06/98 15:15			
A\$3028	17/06/98 15:55			Halocarbons
AS3029	17/06/98 16:15			Halocarbons
AS3030	17/06/98 17:27			Halocarbons
A\$3031	17/06/98 17:47			Halocarbons
AS3032	17/06/98 18:18			Halocarbons
A\$3033	17/06/98 19:14			Halocarbons
AS3034	18/06/98 12:55			Halocarbons
AS3035	18/06/98 13:26			Halocarbons
AS3036	18/06/98 13:57			Halocarbons
AS3037	18/06/98 14:27			Halocarbons
AS3038	18/06/98 14:57			Halocarbons
A\$3039	19/06/98 02:42			Halocarbons
AS3040	19/06/98 04:30			Halocarbons
AS3041	19/06/98 10:11			Halocarbons
AS3041	20/06/98 18:49			Halocarbons
	21/06/98 00:34			Halocarbons
AS3043	21/06/98 02:15			Halocarbons
AS3044				Halocarbons
AS3045	21/06/98 22:27			Halocarbons
AS3046				Halocarbons
AS3047	22/06/98 00:26			Halocarbons
A\$3048				Halocarbons
AS3049				Halocarbons
A\$3050				Hajocarbons
AS3051				Halocarbons
AS3052				Halocarbons
A\$3053	27/06/98 00:00			Halocarbons
A\$3054	27/06/98 21:07			
AS3055	28/06/98 02:29			Halocarbons
AS3056				Halocarbons
AS3057				Halocarbons
AS3026				Halocarbons

CTD 1 - 9/6/98 13:47-14:13, 59°20'N, 19°W, Outside

Lyase	MS				,	Y							
<u>z</u>	GM												
<u>င့</u>	17												
Bact	CPA	7							*				×
200	CS			λ	Y			-					
DMS	RS				<b>&gt;</b>								×
ā	AT												
TCO2	Œ												
Hydr	WB												
Halo	JB												٨
$N_2O$	TF												
$SF_6$	FC		٨			<b>&gt;</b>	Y	Y	¥	X	>	>-	
Sal	JR												
Nuts	TJ	Ă		>-			Y	Ϋ́	Y	Y	¥	Ÿ	Ÿ
AFC	SA												
Phyto	MHL												
HPLC	MHIL												
Chl	MHIL	1									ļ 		
Depth Bottle			14T	8R	12R	101	6T	24T	22T	18T	16T	20R	2T
Depth		0	0	01	101	10	30	45	50	55	100	150	200

CTD2 - 9/6/98, 15:18-15:40, 59°20'N, 19°W, 0-50M Failed!, Outside

	,	_	_		_	_			<u>_</u> _	-,-	- <del>-</del> -	_	_
Lyase	MS												
ž.	GM												
<u></u>	ÐΊ												
Bact	CPA												
Z00	CS												
DMS	RS												
Ĭ	AT	***************************************											
TCO2	JD												
Hydr	WB							-					
Hało	JB												
OzN	TF												
$SF_6$	FC												
Sal	JR.												
Nuts	ŢĴ									Y			
AFC	SA												
Phyto	MHL												
HPLC	MHL												
Chl	MHL												
Depth Bottle		22T	Z4T	18T	20R	18T	14T	12R	10T	8R	T9	4R	2T
Depth		0	0	10	10	25	25	50	50	75	75	100	100

CDT3 - 9/6/98, 16:29-17:05, 59°20'N, 19°W, Outside

Lyase	MS												
N <sub>S1</sub>	ĞM									<b>&gt;</b>	•	<b>&gt;</b>	
14C 15N	FG												-
Bact	CPA					\ \	<b> </b> >-	· >	\ \	>			<b>&gt;</b> -
Z00	S									Ÿ	>		
DMS	RS												
ā	AT												
$TCO_2$	Ωſ	¥				>				Y		7	
Hydr	WB							>-	<b>&gt;</b>	Ϋ́		×	
Halo	JB							Y	<b>*</b>	Ϋ́		<b>&gt;</b>	
N <sub>2</sub> O	TF	X		Ý	Ϋ́	1		×	¥			λ	
$SF_6$	FC												
Sal	JR	¥	Y	Y	Ϋ́	¥	×	Ý	<b>×</b>	Y			
Nuts	TJ	¥		Y									
AFC Phyto	SA							¥	Y	Ϋ́			¥
Phyto	MHL												
HPLC	MHL												
Chi	MHL						Ă	Ϋ́	Å	Y			Ă
Bottle		2	4R	9	8R	10	12R	14	91	18	20R	22	24
Depth Bottle		005	400	300	250	200	150	100	20	10	01	0	0

CTD4 - 10/6/98, 06:01-06:38, 59°N, 22°30°W, Outside

	1		7	<del></del>	1	_	T	т-	<del>-</del>	1	1	1	<del></del>
Lyase	MS	  -								Ϋ́	ı		
14C 15N	ĞM												
14C	IG						***************************************						
Bact	CPA												
Zoo	S								×	>	×		>-
DMS	RS								X	Y	¥		Ϋ́
Q	AT												
TCO2	G.												
Hydr	WB								λ	٨		Ϋ́	>-
Haio	JB							>-	¥	<b>&gt;</b>		¥	λ
N <sub>2</sub> O	TF								-				
$SF_6$	FC												
Sal	JR	Y	<b>~</b>	¥	Ý	Ý	¥	Y	7	<b>≻</b>			
Nuts	ŢĴ	Ϋ́	>	Y	X	Ϋ́	Ϋ́	>	<b>X</b>	<b>&gt;</b>	Ϋ́		Ă
AFC Phyto	SA												
Phyto	MHL												
HPLC	MHL				,								
Chi	MHL						•	Ϋ́	Ϋ́	γ	Y		۲
Bottle		2	4R.	9	8R	10	12R	14	16	18	20R	22	24
Depth		200	400	300	250	700	150	100	50	20	10	10	0

CTD5 - 13/6/98, 07:04-07:55, 59°15'08"N, 20°32'45"W, Outside

Lyase	MS	۸	*												
<sup>14</sup> C <sup>15</sup> N Chlor Lyase	S							;	_	>-		Y		Y	
Z	МЯ											Y			
<u>န</u>	97											Y			
Bact	CPA														
Z00	CS									Y	Y		Υ		
DMS	RS														
Ω	AT						_	_						_	
TCO2	E														
Hydr	WB														
Halo	B														
O <sup>z</sup> N	TF														
SF6	Ů.														
Sal	2		Y	Y	Ă	Y	Υ	Y	>	<b>X</b>	_	>		<u> </u>	-
Nuts	Τ.		Y	Ă	Ϋ́	Ϋ́	Υ	Y	Ă	>	Ϋ́			>	,
AFC Phree	riiytto S.A	5							<b>&gt;</b> -	  -		>	-		7
Phyto	MEI	TATE													
HPLC	MALIE	141112													
Chi	MUI	MITTE				¥	,	X	>	· >		>	-	>	_
Bottle			~	4R	9	8R	10	12R	14	16	2 ~	doc	707 207	777	+7
Depth	•••		200	400	300	250	200	150	100	20	10	2 2	2		>

CTD6 - 13/6/98, 13:05-14:00, 59°30'20"N, 21°07'55"W, Inside

											_					 		_
Lyase	SYV	CIVI			>-													
He	, ca	N.	>- 		>_		7				-						-	
SF6	Ç	اد	>_		<b>&gt;</b>	2	× ;	<u> </u>	×	>	, ;	<u> </u>		>	×		_	
Z.		Σ S																
14 C	,	3														-		
Bact		CPA	>-						×			٨.		<u> </u> ;	χ			
Z00		S	>		<b>&gt;</b>	,-	λ	Y	Y	,	×	>-						
DMS		RS	Y															
a		ΑŢ	Ϋ́		Ϋ́		<b>X</b>	≻	>-	,	,				<b>&gt;</b>			
TCO2		g																
Hydr		WB	¥		<b>&gt;</b>		Y	×	>			_			>-			
Halo		JB	Y		¥		⊹	٨	>	•					Y			
OžN		TF	¥		Υ		>	>-				Y			≻	_		
Sal		Ж																
Nuts	-	TI	<b>&gt;</b>		Υ		>-	>	>	1	<b>&gt;</b>	>	•	X	>-			
AFC	Phyto	SA					<b>&gt;</b>	<b>&gt;</b>		۲ - ا	<b>&gt;</b> -	>	* * *	Y	Y			
Phyto		MHL																
HPLC		MHL									_					-		
Chl		MHL	>	•	×		>	>		Y	<b>&gt;</b>	>	٠ ;	<b>&gt;</b>	<b>&gt;</b>			
Bottle			74T	. a	22T	20R	181	16T		141	10T	£	10	4K	2.T			
Depth			c	>	10		20	25	5	20	88	3	201	150	200			

CTD7 -- 14/6/98, 02:20-02:43, 59°30'19"N, 21°07'47"W

		_		η-	T	-T	Τ-	Į	_		1	Т	-T-	٦
Lyase	MS						>	-						
Z Z	GM	:												
N <sub>51</sub> 2 <sub>71</sub>	LG													
Bact	CPA						>	1		>	T			
Z00	CS												4	ĭ
DMS	RS											>	-	
Ď	AT													
TCO2	JD													
Hydr TCO2	WB													
Halo	JB	>	1				<b>X</b>	Y	Y	7			λ.	
O <sub>2</sub> N	TF													
SF6	FC													
Sai	2													
Nuts	ΤŢ	ì												
AFC	Phyto													<b>,</b>
Phyto	MHI	TATATA												
HPLC	NAGEL	INTERE												
Chl	Mul	INTER	Y	Y	Ă	>~	>	٨	Y					
Bottle			2	4R	9	10	14	16	18	22	20R	12R	24	8R
Depth			200	150	100	80	09	40	20	10	10	10	0	c

CTD 8 - 14/6/98, 03:15-03:28, 59°30'19"N, 21°07'47"W, Inside

			Γ	T	7	_	Τ	1		Ι				Τ				7
Lyase	MS				_					_					+			
Z	GM		>			>	<u> </u>		>			>-		>	٠,		>	
ပ္	TG		>	1		>			>-			>		>	-		>	
Bact	CPA	>-		;	¥				>					,	-		>	
Z00	CS												>	*				
DMS	RS						-											
ā	AT																	
TCO2	Qſ																	
Hydr	WB																	
Hało	JB															L		
OŽN	TF		Ī															
SF6	FC			~														
Sal	IR																	
Nuts	Τ1											,	I		Ÿ			Y
AFC Pr- 42	FHylo S.A	5					Y		,	Ý		,,	ĭ		¥			Y
Phyto	MHI	INTER																
HPLC	Mari	MELL																
Chi	N.C.C.	MILL	Y		,	X		>	-				<u>.</u>	Y				<b>&gt;</b>
Bottle			2	4R		9	8R	Ç	2	12R	1	-	91	18	20R	77.7	22	24
Depth			30	30		70	20		1,	17	-	11		4	4	-	0	0

CTD9 - 14/6/98, 04:20-04:26, 59°30'19"N, 21°07'47"W, Light Profile Only

CTD10 - 15/6/98, 02:27-02:51, 59°28'44"N, 21°04'50"W, Inside

······································	_	F			<del></del> 1		<del></del> -	<del></del>	<del></del> 1		-		
Lyase	MS												
Z.	ВM												
N S	57												
Bact	CPA	<b>X</b>		<b>&gt;</b>						×			
200	S											Y	<u> </u>
DMS	RS			•			Y						
1	AT												
TCO2	Ωſ												
Hydr	WB												
Halo	JB								-				
N <sub>2</sub> O	TF												
$\mathrm{SF}_6$	FC	Y		Ϋ́	Y	Ϋ́	Y	Y	Υ		Ă		
Sai	J.R.	Y	¥	Y	Y	Y	Y	Ϋ́					
Nuts	TJ	γ	Ă	×	>-								
AFC Phyto	SA												
Phyto	MHIL												
HPLC Phyto	MHL												ļ
Chi	MHL	Ϋ́	Ÿ	Ϋ́	ļ								
Bottle		2	4R	9	10	14	16	18	22	12R	24	20R	8R
Depth		200	150	100	80	09	50	30	10	10	0	0	С

CTD11 - 15/6/98, 03:23-03:43, 59°28'44"N, 21°04'50"W, Inside

		36.	A-C	Nuts	Sal	SF6	0. N	Halo	Hydr		DMS	1S	007	Bact	ပ	z.	DMSe	Lyase
			Phyto															
~	IL   MHL	MHL	SA	I	Æ	<u>ت</u>	TE	JB	WB	Qſ	AT	RS	S	CPA	D'I	Μ̈́	PL	MS
~																		
			Y		·							***************************************	-	>-				
-					•											***************************************		
8R Y																		
10 Y																		
12R Y			Ϋ́	À											¥	Ϋ́		
14 Y			٨	÷										Y	Y	Y		
16 Y			Y												¥	Υ		
18 Y			Y											>	>	Y		
20R			Ϋ́															
Z2 Y				Ϋ́		λ									Ϋ́	Y		
24 Y				λ		Ă					*****			Ϋ́	۲	Y		

CTD12 - 15/6/98, 04:08-04:15, 59°28'44"N, 21°04'50"W, Inside, Light Only

CTD13 - 15/6/98, 15:03-15:30, 59°32'50"N, 21°03'34"W, Outside

	_	Τ.	T			<del></del>	1	1	7-	ı	<del></del>		ŧ
Lyase	SW									ŀ			***************************************
DMSe Lyase	PI.												
N <sub>SI</sub>	S												
14C	57												
Bact	CPA												
Z00	CS												
DMS	RS												
Ω	AŢ			X		¥	>	\ \				>	>
TCO2	E												
Hydr	WB			>		Ý	>-	>				<b>&gt;</b>	>
Halo	JB			>-		>-	×	  >-				Ϋ́	>
0 Z 	TF		X	\ <del>\</del>		Ϋ́	<b>~</b>	¥		>-		<b>&gt;</b>	^
SF	FC		λ	Ϋ́		Ÿ	Ϋ́	À		<b>X</b>		Υ	>
Sal	Æ												
Nuts	Ω												
AFC Phyto	SA												
Phyto	MHL												
HPLC	MHL												
Chl	MHL		<b>&gt;</b>					Ϋ́					
Depth Bottle		12R	16T	18T	20R	22T	24T	2T	4R	19	8R	10T	14T
Depth		10	10	10	10	10	10	100	100	100	100	100	100

CTD14 - 16/6/98, 02:15-02:35, 59°33'59"N, 21°06'54"W, Outside Upwelling Region

-			,		_	_								
I vace	Agg Ca	SMS												
Ήe	}		Z							<b> </b>				>
DMSe		Ы								-				
N <sub>S</sub> I	:	M.P.												
N <sub>51</sub> 15N	)	IG												
Bact		Г	>		<b>&gt;</b>									
Zoo	}	CS										\ \	\ \	
S	}	RS												
DMS		AT												
TCO,	•	13												
Hvdr TCO,		WB												
Halo		JB			<u>}</u>									
N <sub>2</sub> O		TF												
SF6		FC	×		X			Ϋ́	<u> </u>	Ϋ́				Y
Sal		JR	<b>~</b>	>	<b>*</b>	\ \	٨	Υ	Ϋ́	¥				
Nuts		TJ	Ϋ́	<b>&gt;</b>	λ	<u> </u>	Ϋ́	٨	Υ		7		¥	
<b> </b>	Phyto	SA			\ \									
Phyto		MHL				ļ			<u>.</u> .				<u>                                     </u>	
HPLC		MHL		-	L				<u> </u>					
ChI		MHL											-	
		L	2	48	9	01	14	16	18	22	12R	20R	8R.	24
Depth Bottle			200		100	50	25	20		10	10	0	0	0
L			_	_	يـــــــــــــــــــــــــــــــــــــ								_	

CTD15 - 16/6/98, 03:05-03:20, 59°33'59"N, 21°06'54"W, Outside Upwelling Region

		_			_	_							1
Lyase	MS												
DMSe Lyase	PL												
Ν̈́ςι	GM				>-	×	Y	Ϋ́	Y		>		
14C	97				Y	Y	Ϋ́	Å	Ϋ́		<b>&gt;</b>		
Bact	CPA		>										
Zoo	CS								Ÿ	Y			
DMS	RS												
Ω	AT	>		<b>~</b>		Ă						٨	
TCO <sub>2</sub>	Ü												
Hydr	WB	7		¥						<b>*</b>		Y	
Halo	JB	X		λ		>-						¥	
N <sub>2</sub> O	TF	Y		>-		\ <del>`</del>						Ϋ́	
SF <sub>6</sub>	FC										<b>&gt;</b>		
Sal	JR												
Nuts	ŢĴ												
AFC Phyto	SA	Ý					>	\\ \		7	γ		
Phyto	MHL	¥											
HPLC	MHL												
CPI	MHL	λ	>	٨	>	· >-	· >	\ \ \		<u>\</u>		Ϋ́	
Depth Bottle		2	4R	9	8R	9	128	14	16	18	20	22	
Depth		9	50	40	30	2 02	17	1	4	4	0	0	

CTD16 - 16/6/98, 04:01-04:09, 03:05-03:20, 59°33'59"N, 21°06'54"W, Outside Upwelling Region

							_		_	_
Lyase	MS	۲ ;	٠,							
DMSe Lyase	PL		***************************************							
N <sub>S1</sub>	GM									
Bact <sup>14</sup> C	FG									
Bact	CPA									
Z00	CS	Î								
DMS	RS	×	>-				ļ			
D	ΑT									
TCO2	JD									
Hydr	WB									
Halo	JB									
N <sub>2</sub> O	TF							-		
SF <sub>6</sub>	FC									
Sai	JR									
Nuts	TJ							***************************************		
AFC Phyto	SA									
HPLC Phyto	MHL			-			***************************************			
HPLC	MHL									
Chl	MHL									
Depth Bottle		2	4R							
Depth		4	4							

CTD17 - 17/6/98, 02:14-02:35, 59°33'45"N, 20°46'07"W, Inside

, s	T	T	T	T		T	T	Τ	T	Τ	Τ	Τ	
Lyase	We	CVA	+	-	-	-	_	-	_		1	_	
He	Z						-		>	-		>	
DMSe	ρί	7											
N S	SM S												
14C	<u>ر</u>												
Bact	CPA	>			>	,				>			Y
Z00	S	3									>	,	¥
DMS	SS												>
Ô	AT					-			-		-		
TCO2	ij												
Hydr	WB												
Halo	JB							-					
N <sub>2</sub> O	TF												
$SF_6$	5	>		×		λ	>-	X	<b>X</b>			¥	
Sal	Æ	<u>}</u>	X	Ÿ	>-	Y	Y	7		λ			
Nuts	II	X	<b>&gt;</b>	Ϋ́	Υ	Y	7	>		λ.			¥
AFC Phyto	SA												
Phyto	MHL												
HPLC	MHL												
Chl	MHL	Ϋ́		Υ	Ÿ	Ϋ́	Ϋ́	Y					
Bottle		2	4K	9	10	14	16	18	22	12R	20R	24	8R
Depth		200	150	100	35	30	25	20	10	10	0	0	0

CTD18 - 17/6/98, 03:09-03:24, 59°33'45"N, 20°46'07"W, Inside

_	1	_				_	_		<del></del>	_	1	_	
He	NA	; >	-							^	4		
DMSe	īd												
Z <sub>SI</sub>	35			>	1 >	\ \ \	· >	\\.\.\>	,		¥		
٦ <sub>+</sub> 1	<u>ر</u>			>	· >	· >	\ \>	\  >		>			
Bact	CPA	>		٨	· >	1	>	. >			X		
Z00	S								>	*			
DMS	RS												
a	AT												
TCO <sub>2</sub>	2												
Hydr	WB												
Halo	ar Br								-				
O <sub>2</sub> N	TF												
SF6	FC												
Sal	Ж												
Nuts	Ţ												
AFC Phyto	SA	≻		<u>\</u>	Υ		¥	Υ		Ϋ́			
Phyto	MHL												
HPLC	MHL												
CFI	MHL	¥	>-		٠			¥			>-		
Depth Bottle		2	4R	9	8R	10	12R	14	16	18	20R		
Depth		50	40	30	20	17	11	4	4	0	0		

CTD19 - 176/98, 04:15-04:22, 59°33'45"N, 20°46'07"W, Inside, Light Only

CTD20 - 18/6/98, 03:03-03:22, 59°31'26"N, 21°00'01"W, Inside

1	Т	1	T		····T	Т	Т	1	Т		7	<del></del>	-
Lyase	MS												
- 1	Z.										≻	X	
DMSe	PL												
<sup>14</sup> C L <sup>1</sup> N DMSe	₩ B												
္	27												
Bact	CPA			×	Y					Y			
Z00	S												
DMS	RS												
Ω —	AT					Υ		Y			<b>&gt;</b>	Ϋ́	
TCO2	Ωſ												
Hydr	WB							Ý			>	Ϋ́	
Halo	JB												
N <sub>2</sub> O Halo	TF	<b>&gt;</b>		Υ		Y			Υ		⋆	Ϋ́	
SF <sub>6</sub>	FC	Y		Υ		λ		Y	Y		Ÿ	Ϋ́	
Sal	JR	Y	Y	Y	Ϋ́	Ÿ		Y					
Nuts	TJ												_
AFC Phyto	SA							Å		Ă			^
Phyto	MHL												
ЭЛАН	MHL												
Chl	MHL												
Bottle		2	4R	9	8R	10	14	16	18	20R	22	24	125
Depth		200	150	100	50	30	.25	20	15	10	10	0	

CTD21 -- 18/6/98, 04:11-04:24, 59°31'26"N, 21°00'01"W, Inside

										_			
Lyase		MS								Ϋ́	Υ		
He		NA.											
DMSe	200	PL											
Z 2		ĞM											
r4C		57											
Bact		CPA	>	><		Ý	¥			Y			
Z00		CS											
DMS		RS											
ìa		AT		×	***************************************								
TCO2		Ω											
Hydr		WB		٨									
Halo	***************************************	JB											
O²N		TF		≻									
$SF_6$		FC								<b>&gt;</b> -			
Sal		Æ											
Nuts		Ω			Å		×		¥			 ***************************************	*****
AFC	Phyto	SA											
Phyto	- 1	MHL											
ЭЛАН		MHL											
Chi		MHL	¥	<b>&gt;</b>	٨	Ϋ́	×	X		>			
Depth Bottle			2	9	10	14	16	81	22	24	20R		
Depth			100	50				4	4				

CTD22 - 19/6/98, 02:39-02:46, 59°31'26"N, 21°00'01"W, Inside

Lyase	MS			Ÿ					
He	No.								
DMSe He	PL								
$N_{SI}$	GM								
J⁴C	PT								
Bact	CPA		<b>&gt;</b>						
Z00	8	>	· >-						
DMS	RS			Ϋ́					
ā	AT					-			
TCO <sub>2</sub>	Œ								
Hydr	WB								
Halo	JB		<b>*</b>			-			
OzN	TF								
$SF_6$	FC			Ϋ́					
Sal	H.								
Nuts	TJ		Y						
AFC Phyto	SA			Y					
	MHL		27						
HPLC	MHL		Ϋ́						
СЫ	MHL		Y						
Depth Bottle		4R	2	9					
Depth		0	0	0					

CTD23 - 20/6/98, 22:09-22:33, 59°33'12"N, 21°01'36"W, Inside

Lyase	MS							-				
He	Z							>	· >	٠		
DMSe	PI.	1										-
Z <sub>SI</sub>	W.S											***************************************
N <sub>SI</sub>	97 179						-					
Bact	CPA											
Z00	బ											
DMS	RS		-									
Ĭ	AT								Y			
TCO2	£											
Hydr	WB								Y			
Halo	JB								Ϋ́			
N <sub>2</sub> O	TF							>	×			
SF6	FC	¥	Ϋ́	Y	¥	Υ	Y	X	<b>&gt;</b> -			
Sal	J.R											
Nuts	ŢJ								⊁			
AFC Phyto	SA								Ă			
Phyto	MHL											
	MHL								<b>→</b>			
Chi	MHL								۲			
Bottle		2	9	10	14	16	81	22	24	20R	12R	
Depth		100	80	70	65	09	20	10	4	4	4	

CTD24 - 21/6/98, 03:02-03:20, 59°33'12"N, 21°01'36"W, Inside

Depth         Bottle         Chl         HPLC         Phyto         AFC         Nus         Sal         Fig         Hod         FC         TT         Halo         Hydr         TCOs         DMS         GS         CPA         IG         FC         TT         Halo         Hydr         TCOs         CPA         IG         FC         TT         Halo         Hydr         TCOs         CPA         IG         FC         TCOs         TCOs         CPA         IG         PC         PC <t< th=""><th></th><th></th><th></th><th>_</th><th>_</th><th>_</th><th><del></del></th><th></th><th></th><th></th><th></th><th></th><th></th><th></th></t<>				_	_	_	<del></del>							
Bottle         Chl         HPLC         Phyto         AFC         Nuts         Sal         SFe         Halo         Hydr         TCO2         DMS         200         Bact         HG         IS         IS         DMS           2         Y </td <td>Lyase</td> <td>Mg</td> <td>OT.</td> <td></td>	Lyase	Mg	OT.											
Bottle         Chi         HPLC         Phyto         AFC         Nuts         Sal         SF6         NgO         Halo         Hydr         TCO <sub>2</sub> DMS         Zoo         Bact         H <sub>C</sub> IS           2         Y	He	Nd												
Bottle         Chi         HPLC         Phyto         AFC         Nuts         Sal         Sfe         N2O         Halo         Hydr         TCO2         DMS         200         Bact           2         MHL         MHL         SA         TJ         JR         FC         TF         JB         WB         JD         AT         RS         CS         CPA           4R         Y	DMSe	Ĭd												
Bottle         Chi         HPLC         Phyto         AFC         Nuts         Sal         Sfe         N2O         Halo         Hydr         TCO2         DMS         200         Bact           2         MHL         MHL         SA         TJ         JR         FC         TF         JB         WB         JD         AT         RS         CS         CPA           4R         Y	Ž.	GM												
Bottle         Chl         HPLC         Phyto         AFC         Nuts         Sal         SF6         N2O         Halo         Hydr         TCO2         DMS         DMS         Zoo           2         Y         Y         Y         Y         Y         Y         X	14C	ر ا	2											
Bottle         Chl         HPLC         Phyto         AFC         Nuts         Sa         SF6         N2O         Halo         Hydr         TCO2         TMS           2         Y         Y         Y         Y         X	Bact	CPA	>		<b>&gt;</b>			>			<u> </u>			Ϋ́
Bottle         Chl         HPLC         Phyto         AFC         Nuts         Sal         SF <sub>6</sub> N <sub>2</sub> O         Halo         Hydr         TCO <sub>2</sub> DM           2         Y	Z00	S			λ								Ϋ́	
Bottle         Chl         HPLC         Phyto         AFC         Nuts         Sal         SF6         N2O         Halo         Hydr         TCO2           2         Y	MS	RS										<u>\</u>		
Bottle         Chl         HPLC         Phyto         AFC         Nuts         Sal         SF6         N2O         Halo         Hydr           2         Y         Y         Y         Y         WB           4R         Y         Y         Y         Y         WB           6         Y         Y         Y         Y         Y           10         Y         Y         Y         Y         Y           14         Y         Y         Y         Y         Y           16         Y         Y         Y         Y         Y           18         Y         Y         Y         Y         Y           22         Y         Y         Y         Y         Y           20         Y         Y         Y         Y         Y           22         Y         Y         Y         Y         Y           24         X         Y         Y         Y         Y           24         X         Y         Y         Y         Y           24         X         Y         Y         Y         Y           24         X<	Ω	AT		1	>	>	·   >-	>						
Bottle         Chl         HPLC         Phyto         AFC         Nuts         Sal         SF <sub>6</sub> N <sub>2</sub> O         Halo           2         Y         Y         Y         Y         Y         N         Y         N		J.									-			
Bottle         Chi         HPLC         Phyto         AFC         Nuts         Sal         SF <sub>6</sub> N <sub>2</sub> O           2         Y         T         Y         Y         TF           4R         Y         Y         Y         Y         Y           10         Y         Y         Y         Y         Y           14         Y         X         Y         Y         Y           16         Y         Y         Y         Y         Y           18         X         X         Y         X         X           20         X         X         X         X         X           24         X         X         X <td>Hydr</td> <td>WB</td> <td></td>	Hydr	WB												
Bottle         Chi         HPLC         Phyto         AFC         Nuts         Sal         SF <sub>6</sub> N <sub>2</sub> O           2         Y         T         Y         Y         TF           4R         Y         Y         Y         Y         Y           10         Y         Y         Y         Y         Y           14         Y         X         Y         Y         Y           16         Y         Y         Y         Y         Y           18         X         X         Y         X         X           20         X         X         X         X         X           24         X         X         X <td>Halo</td> <td>JB</td> <td></td> <td></td> <td>X</td> <td></td> <td>X</td> <td>&gt;-</td> <td></td> <td>&gt;-</td> <td></td> <td></td> <td></td> <td></td>	Halo	JB			X		X	>-		>-				
Bottle         Chil         HPLC         Phyto         AFC         Nuts         Sal           2         Y         TJ         Y <td></td> <td>TF</td> <td></td> <td></td> <td>&gt;-</td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td> <td></td>		TF			>-									
Bottle         Chl         HPLC         Phyto         Nuts           2         Y         TJ           4R         Y         R         TJ           10         Y         R         R           14         Y         R         R           16         R         R         R           22         R         R         R           24         R         R         R           12R         R         R         R           24         R         R         R           3R         R         R         R	SF <sub>6</sub>	FC	¥		λ	>-	Å	Ϋ́	×	>		>-		
Bottle         Chl         HPLC         Phyto         AFC           2         Y         Phyto         Phyto           4R         Y         R         Phyto           6         Y         R         Phyto           10         Y         R         P           16         Y         R         P           16         Y         R         P           18         Y         R         P           22         Z         R         P           24         R         R         P           12R         R         R         P           12R         R         R         P           12R         R         R         P	Sal	JR	×	>-	٨	>	>-	Ϋ́	Ϋ́			>		
Bottle         Chi         HPLC         Phyto           2         Y         MHL         MHL           4R         Y         R         P           6         Y         R         P           10         Y         R         P           14         Y         R         R           16         Y         R         R           22         R         R         R           24         R         R         R           12R         R         R         R           12R         R         R         R	Nuts	TJ												
Bottle         Chl         HPLC           2         Y         HRL           4R         Y         6           10         Y         11           16         Y         12           18         Y         12           22         20R         24           24         12R         8R	AFC Phyto	SA												
Bottle Chi  2 Y 4R Y 6 Y 10 Y 14 Y 16 X 16 X 22 Z 24 Z 24 Z 8R	Phyto	MHL												
Bottle 2 2 4R 6 10 10 14 14 18 22 22 20R 24 12R	HPLC	MHL												
	Chl	MHL	≻	>-	Ϋ́	٨	<u> </u>							
Depth 200 150 100 80 60 40 20 10 10 0 0	Bottle		2	4R	9	1.0	14	91	81	22	20R	24	12R	8R
	Depth		200	150	100	80	09	40	20	01	10	0	0	0

CTD25-21/6/98, 03:47-03:59, 59°33'12"N, 21°01'36"W, Inside

Lyase	MS	Crar						٨			-		
He	PN				-						-	-	
DMSe He	PI												
N <sub>SI</sub> D <sub>FI</sub>	S.	>	· >-	· >-		\ \ \					Y		
14C	1,G	<u> </u>	<u>\</u>	<u> </u> >-	·   >-	· >-		<u>\</u>			>		
Bact	CPA												
00Z	SS	}					***************************************		٨	>-			
DMS	RS	<b>*</b>	>		<u> </u>			Ϋ́					
Ω	AT	<b>&gt;</b>	>		>-		Ă		Ϋ́				
TCO2	ξ		Y		>		٨						
Hydr	WB		γ		>-		×		Y				
N <sub>2</sub> O Halo	JB												
N <sub>2</sub> O	TF												
SF <sub>6</sub>	FC								⊁				
Sal	R												
Nuts	II.	Å			>	Y					Ϋ́		
AFC Phyto	SA	<b>&gt;</b>	Y	Ϋ́	Ϋ́	×					Ϋ́		
Phyto	MHL										32		
HPLC	MHL												
Chi	MHL	٨	Y		Ÿ	Ϋ́					Ă		
Bottle		2	9	10	14	16	18	22	24	4R	8R		
Depth		30	20	17	11	4	4	0	0	0	0		

CTD26 - 21/6/98, 15:05-15:33, 59°34'47"N, 21°08'38"W, Edge

_													-		
Fe	1.1	CY.	>	1	>	-			٨	,			>	\ \	
He	Nd	111													-
DMSe	Ιd													<b>\</b>	
Z <sub>51</sub>	GM	,													1
ا <del>ا</del> د	1.6	2													
Bact	CPA														,
Zoo	8														
DMS	RS														
ã	AT			>			<b>&gt;</b>			<b>~</b>	>	>		×	
TCO2		  -		<b>&gt;</b>							>-				
Hydr	WB	λ.					>-			<b>}</b>	Ϋ́	>			***************************************
Halo	38														
O <sup>z</sup> N	TF														
$SF_6$	F.	>-		<b>}</b>		>-	<b>×</b>	¥		Ϋ́	Y	X			I
Sal	JR														
Nuts	TI	>	ļ	Ϋ́		<b>}</b>	7	<b>*</b>		Υ	λ	<b>Y</b>		×	
AFC Phyto	SA														
Phyto	MHL	40		39		38	37	36		35	34	33		41	
HPLC	MHL	×		<b>X</b>		Y	X	<b>≻</b>		٠.	Y	×		Ý	
Chl	MHL	<b>}</b>		¥		×	Ÿ	<b>&gt;</b>		Ϋ́	Ϋ́	Ϋ́		Y	
Bottle		2T	4R	6T	8R	T01	14T	16T	20R	18T	22T	24T	12R	10L	
Depth		150	150	80	80	70	09	45	45	40	25	0	0	0	

CTD27 - 22/6/98, 12:00-13:00, 59°53'N, 21°15'23"W, Edge

	_	_											
Lyase	Me	>	*		>		>	-			٨	1	
He	Na		-				>	¥	>	·   }			
DMSe	ρĩ											>	
$N^{21}$	ME												
ر بر	1.6	)											
Bact	CPA												
Z00	S										Ϋ́		
DMS	RS												
ď	AŢ	>		Ϋ́	Y	>	\ \		\ \	\ \		Υ	
TCO2	Q.						-						
Hydr	WB	¥		Y	λ	>	X		X	Y			
Halo	J.B	λ			×	Ă	Y		Y	Ϋ́			
OŽN	TF												
$SF_6$		×	<b>}</b>	*	Υ	Ϋ́	¥		Ϋ́	<b>}</b>			
Sal	Æ												
Nuts	ΙΊ	7	¥	Ϋ́	¥	Ÿ	<u>۲</u>		\ \		Ϋ́	Ϋ́	
AFC Phyto	SA												
Phyto	MHL	. 55		54	53	52	51	50			46	48	
HPLC	MHL											Ϋ́	
Chi	MHL	Ÿ		<b>&gt;</b>	Ÿ	Ϋ́	Ÿ	<b>&gt;</b>			Y	λ	
Bottle		2T	6T	10T	14T	16T	18T	20R	22T	24T	4R	10L	
Depth		100	80	70	9	50	25	10	10	0	0	0	

CTD28 - 23/6/98, 02:53-02:59, 59°54'19"N, 20°44'30"W, Inside

He	N.								
DMSe He	PL								
N <sub>C</sub> ISN	GM								
14C	ΓG								
Bact	CPA			Y					
Zoo	CS	Y	Y						
DMS	RS								
Ğ	AT								
TCO2	UC	λ							
Hydr	WB								
Halo	JB								
N <sub>2</sub> O	TF								
$SF_6$	FC								
Sal	JR								
Nuts	ŢĴ							1	
AFC Phyto	SA								
Phyto	MHL								
HPLC	MHL	X							
СЫ	MHL		_						
Depth Bottle		2	4R	9					
Depth		0	0	0					

CTD29 - 23/6/98, 03:22-03:42, 59°54'19"N, 20°44'30"W, Inside

		Т	Т	-	1	T		Т	Т				٦
Lyase	MS										¥	Υ	Y
He	N.	1											
DMSe	PĽ												
N <sub>SI</sub>	GM								Y				Y
$\mathcal{D}_{Pl}$	FG				;		Ϋ́		Ϋ́	¥	Y		Y
Bact	CPA	¥	Y			¥		Y	Y	Y			Y
Z00	CS											Y	
DMS	RS										7		
ā	AT			Y	¥	Ϋ́		Y	Y	Ϋ́	≻		
TCO2	JD			Y	Υ								Y
Hydr	WB				Ă	¥		Ϋ́	Ý	Y		•	
Halo	JB			Ý	Y	¥		<b>&gt;</b>	٨	Y			
O <sup>2</sup> N	TF	Ý		Ý	<b>&gt;</b>	<b>&gt;</b>		Υ	Ϋ́	Ϋ́	Ϋ́		
SF6	FC	Y		Ă	¥	7		<b>~</b>	×	×			
Sai	JR	Ϋ́											>
Nuts	TJ	Y	Å		<b>&gt;</b>	<b>&gt;</b>		¥	٨	>		>-	>-
AFC Phyto	SA			¥		<b>}</b>	λ		¥	¥			¥
Phyto	MHL		99		65	49		63	62	61	09		59
HPLC	MHL		¥		×	×		×	Ϋ́			>-	×
Chi	MHL	٨	Y		>	<b>&gt;</b>		Y	Ý	Ϋ́	>		¥
Bottle		2	4R	9	10	14	12R	91	18	22	24	20R	8R
Depth		200	100	100	09	40	30	30	20	10	4	4	0

CTD30 - 23/6/98, 05:58-06:17, 59°54'19"N, 20°44'30"W, Inside

Γ	Ţ	т	T	т-	Т	T-	т-	1	1	Τ	<del></del>		_
DMSe	PI												
Z Z	ĞM												
14C	2											>-	
Bact	CPA		Y		×	Y	*	X				¥	
Zoo	S												
DMS	RS										Ϋ́		
ប្រ	AT				Ϋ́	۲	>-	7			λ		
TCO2	100												
Hydr	WB			-							<b>&gt;</b>		
Halo	JB						¥	<b>&gt;</b>			<u> </u>		
O <sup>z</sup> N	TF		¥	Y		×	>	Ϋ́		Y	Ϋ́		
$SF_6$	F.		X	Y	Ϋ́	<b>}</b>	Ϋ́	¥		Ϋ́	×		
Sal	JR	>-									¥		***
Nuts	13	Y	>-	¥	≻	Ϋ́	Y	¥	Ϋ́		Ϋ́		
AFC Phyto	SA	¥	<b>&gt;</b> -	<i>&gt;</i>	×	¥	Y	Y	Ă		Y		
Phyto	MHL			73	72	71	70	69		89	29		
нес	MHL				Y	Υ	Y	Y		, Y	λ		
СЫ	MHL		Y	Υ	Ă	Υ	Y	Y		Ϋ́	Y		
Bottle		4R	2	9	10	14	16	18	20R	22	24	12R	
Depth		200	100	09	40	30	20	10	4	4	0	0	

CTD31 - 23/6/98, 08:05-08:20, 59°52'03"N, 20°34'17"W, Edge

DMSe	<u> </u>				÷				
<sup>14</sup> C <sup>15</sup> N DMSe	МS		λ	λ					
14C	TG						-		
Bact	CPA								
002	S	-							
DMS	RS					***************************************			
Q	AT								
$TCO_2$	U								
Hydr	WB								
Halo	JB			>					
N <sub>2</sub> O	TF			¥					
$\mathrm{SF}_6$	F.								
Sal	Ħ,	7		7					
Nuts	TJ		<b>&gt;</b> -	Ϋ́					
AFC Phyto	SA								
Phyto	MHL							-	
HPLC Phyto	MHL								
Chi	MHL								
Depth Bottle		2	4R	9					
Depth		200	20	0					

CTD32 - 23/6/98, 10:23-10:43, 59°52'03"N, 20°34'17"W, Edge

	-	_	-1	_	_	-[			Т			<del></del>	 1	7	
Lyase	MS	-							>	ĭ					
<sup>14</sup> C <sup>15</sup> N DMSe Lyase	PL														
N <sub>S1</sub>	GM														
$^{14}\mathrm{C}$	FG								;	,					
Bact	CPA		Y		>	⊁	>-	۸		٨					
Z00	CS														
DMS	RS									X					
ā	AT				Y	<b>&gt;</b> -	<b>*</b>	>	ĭ	>					
TCO <sub>2</sub>	Ωſ														
Hydr TCO <sub>2</sub>	WB									χ				,	
Halo	JB					Y	>	`		>-					-
N <sub>2</sub> O	TF							>	ĭ	>					
SF6	FC	Y	Y	Y	Ϋ́	>-	>	, ,	,	<b>&gt;</b>			-		
Sal	JR	Y								>-					-
Nuts	TJ									>-					
AFC	SA	ļ													
Phyto	MIHIL														
HPLC	MHL														
망	MHL									<b>&gt;</b>	-				
Depth Bottle		2	9	01	14	1,4	2 9	81	22	24					
Depth		500	100	9	40	30	200	07	10	c					

CTD33 - 23/6/98, 13:11-13:35, 59°50°52", 20°39°35"W, Edge/Outside

	<del></del>	,	_		_	_			<del></del>	<del></del>			_	 	_	ſ	1
DMSe	),G	7,									>				***************************************		
<u>z</u>	2	i divi															
N <sub>51</sub> D <sub>b1</sub>	0,1	אַן															
Bact	740	CFA	1	7	>	>	<b>&gt;</b> -	***************************************	χ	Y	>-						
Zoo	20	3		Y	Y	X	<b>&gt;</b>		<b>&gt;</b>	Y	>-						
4S	5	2															
DMS		Al		>	Υ		>-		>	Y							
TCO,	4	Ωſ									Ϋ́						
Hydr		WB		Y	>-		<b>}</b>		Y	<b>&gt;</b> -	Ÿ						
Halo		JB				Y	Ϋ́		Y	Ă	>-						
N <sub>2</sub> O		ŢF		<b>≻</b>	¥		¥		<u> </u>		Ϋ́	•					
$SF_6$		FC	Y	Y	>									 <b>-</b>			
Sai		JR	Y								Y						_
Nuts		Ţĵ	Y	Y	<b>&gt;</b>	<b>×</b>	Υ		<b>&gt;</b> -	>	>						
AFC	Phyto	SA		¥	  >-	Y	Ϋ́		Υ	>	>						
Phyto		MHL	82	18	08	62	78		77	9/	75						
HPLC		MHL	¥	\ \	>	×	¥	•	Ϋ́	٨	\ \ >	•		T			_
Chi		MHL	×	>	  >	Ϋ́	<b>&gt;</b>		\ \	>	. >	•					
Bottle	لىــــل	<u> </u>	2T	6T 4R	10T	14T	16T	12R	18T	27.T	24T	20R	8R				
Depth			200	100	9	40	30		20	10	2, 0	>					

CTD34 - 26/6/98, 21:05-21:30, 61°11'50"N, 20°31'14"W, Outside

DMSe	PL										
<u>%</u>	ĞM									 	
14C	27							***************************************			
Bact	CPA										
Zoo	S								<b>&gt;</b> -		 ļ 
DMS	RS										
Ω	AT	<b>&gt;</b> -	>	>	≻	¥	<b>&gt;</b>				
TCO2	30		>	٨					X		
Hydr	WB		Y	X		Y	Ă				
Haio	ar	Ă									
N <sub>2</sub> O	TF		<b>&gt;</b>	<b>&gt;</b> -		>-	×	Y	Y		
$SF_6$	FC										
Sal	JR	:									
Nuts	] TJ	٨	<b>&gt;</b>	Ϋ́	٨	X.	Υ	Ϋ́	λ		
AFC Phyto	SA										
Phyto	MHL	101	102	103	104	105	901				
HPLC	MHL	¥	λ	Y	Ă	Ϋ́	λ			 	
Chl	MHL	¥	Χ	Ϋ́	À	Ϋ́	Ă	۲	٨		
Bottle		24T	22T	18T	16T	14T	10T	L9	2T		
Depth		0	7	20	30	40	09	100	700		

CTD35 - 27/6/98, 02.18:02.41, 60°25'54"N, 20°39'37"W, Inside Upwelling Region

			_	Ę	_	ł		<b></b>		T	1	_
Lyase	MS										X	
DMSe Lyase	PĽ											
N.	GM											
ا <del>ب</del> ر	FG							Ÿ				
Bact	CPA	¥	Y		Y							
Zoo	CS						Ϋ́	Ϋ́				 
DMS	RS											
Ď	AT			X	<b>,</b>	×						
TCO2	ar											
Hydr	WB		¥	≻	Y							
Halo	JB		>-	>	Y							
O <sup>z</sup> N	TF	Ϋ́	>	>-	×							
SF <sub>6</sub>	FC											
Sal	JR	¥									Y	
Nuts	IJ	Ÿ		X								
AFC Phyto	SA											
Phyto	MHL	128		108								
HPLC Phyto	MHL	Ă		Ϋ́								
СЫ	MHL	Å		¥								
Depth Bottle		2	9	10	14	91	4R	18	22	24	8R	
Depth		200	100	09	40	01	4	4	0	0	0	

CTD36 -- 27/6/98, 03:25-03:45, 60°25'54"N, 20°39'37"W, Inside Upwelling Region

		_				1	- 1				- 1			γ-			3
DMSe	PL																
<u>2</u>	ВМ	>-	Y		۸	, ;	Y		**	×							
N <sub>51</sub>	1.G	>-	¥		>	1;	X		;	X		>-					
Bact	CPA	٨		>-		,	Y					>					
Zoo	CS										>-						
DMS	RS																
ได	AT		>					<b>&gt;</b>		Y		Ϋ́		-			
$TCO_2$	Ωſ																
Hydr	WB		<b>X</b>					<b>&gt;</b>		<b>&gt;</b>		Y					
Halo	JB				-			>	-	>-		>				-	
N <sub>2</sub> O	TF		λ					>		<b>&gt;</b> -		>					
$SF_6$	FC									~							
Sal	JR																
Nuts	TJ		Ϋ́					>	,	<b>&gt;</b>							
AFC	SA	<b>*</b>	•	>	7	Y	À			X		٨	*				
Phyto	MHL	132	131					120	130		129	127	777				
HPLC	MHL	>	• >						I		>	· >	1				
Chi	MHL	>	, A					<u> </u>	X		>	· >	-		ļ		~
Bottle		ΔR	¥ 4	ao	A6	10	128	121	<del>*</del>	16	18	2,7	<del>+</del> 7				
Depth		30	200	25	07	17	1-		<u></u>	4	4						

CTD37 - 27/6/98, 21:30-21:45, 59°45'38"N, 21°W, Inside

			_	_	_	-,			_			,	_	_,
Lyase	MS	2	- ;	×	,	X		>	×.	≻				
DMSe	PL													
Z	GM											-		
14C	FG													
Bact	CPA	;	<u> </u>				¥	Y	<b>~</b>		>			
Z00	cs													
DMS	RS		-											
<u>α</u>	AT									>				
$TCO_2$	Ωſ		>-		Υ			×	>	<b>&gt;</b>				
Hydr	WB		j							>	Č.			
Halo	JB				Ϋ́	¥	Υ	≻	>	· >				
O <sub>z</sub> N	TF		>		Υ		×	>-	>	· >				
SF <sub>6</sub>	FC		٨	X	<b>&gt;</b> -	Y	<b>≻</b>	>	>	\ \>	¥			
Sal	JR													
Nuts	ŢĴ		٨	Y		λ		>	>	• >	-			
AFC Phyto	SA													
Phyto	MHL		138	137		136		135	127	123	155			
HPLC	MHL		Ă	⊁		>		>		- >	ĭ			
Chl	MHL		Y	Ϋ́		<b>X</b>		>	<b>\</b>	× 3	ĭ			
Bottle		4R	2T	6T	101	14T	16T	187	101	177	1 57	SR.		
Depth		200	100	70	50	40	30	3 6	07	≧ ,	7	2		

CTD38 - 28/6/98, 02:09-02:33, 59°48'26"N, 20°57'33"W, Inside

DMSe		71		T										
N <sub>51</sub>	+	<u> </u>	-				_							
14C		+												1
Bact	Yay	<u>ا</u>	*	>	-	1	;	٦				4		^
Z00	3	3										>	-	1
DMS	20	3												>
ā	ΔT	3			>	1	>	ĭ		\ \ \	.   >	,		>
TCO2	E			>	< >	-		-				-		
Hydr	WR	2			>	•	>	I			>			Λ
Halo	Œ				>		>	ī		<u>\</u>	<b>\</b>			
N <sub>2</sub> O	ŢŢ				>	į				>-		-		
$SF_6$	<u> </u>			>	\ \	· >	. >	,	Y	Ϋ́	>			>
Sal	Æ	>	,										<u></u>	
Nuts	TJ			<u> </u>	<u></u>		٨	,		<b>≻</b>	*		\ <u>\</u>	>
AFC Phyto	SA			>	X					>			>	
	MHL				147					145	144		143	
HPLC	MHL				<b>&gt;</b>					Y			×	
<del>-</del> 5	MHL			>	\ <u>\</u>		Υ			Y	Y		X	٨
Bottle		4R	8R	9	10	14	16	10	01	22	2	12R	20R	24
Depth		200	150	100	70	09	50	40	7	30	10	4	4	0

CTD39 - 28/6/98, 03:18-03:41, 59°48'26"N, 20°57'33"W, Inside

r	_	_	_			7	_		γ-	· -		_	- 7	_		_	_		
Chl		2	3	χ.								)	_	-			;	Y	
DMSe		id																	
N Si		چ			<b>~</b>		1		>	1		>	۲		X				>
14°C		<u></u>			>-	>	ı		Λ	-		^	*	;	X				<b>&gt;</b>
Bact		CPA																	
Z00		S											,	×		>			
DMS		RS		1															
Ω		ΑŢ					-						^	-					
TCO2		2																	
Hydr		WB											-						
Halo		38				_													
N <sub>2</sub> O	1	<u>-</u>																	
$SF_6$	C L	FC				•••		-											
Sal	5	JK																	
Nuts	,	13					>	,								_			
AFC Phydo	2,4,3,6	9A	Y	>	T I	<b>&gt;</b>					,	<u></u>		٨				,	1
Phyto	Mari	INITITY	146				148					144					142		
HPLC	Mur	IVILLE	>		;	Y					>	¥					<b>&gt;</b>		
F	MILI	יייני	<b>≻</b>				<b>&gt;</b> -												
Depth Bottle			2	9	,	10	14		61	18	doc	ZUK	22	8R	128	177	24	٩V	AT-
Depth			50	30	200	70	20	1.2	/1	11	1.1	1 1	4	4	4	-	0	C	

CTD40 - 28/6/98, 14:19-15:02, 59°49'57"N, 20°41'47"W, Edge/Outside

Š	T					7			ŀ				
DMSe	겊			_	-	>			>	٨	<b>&gt;</b>	>	<b>&gt;</b>
N <sub>2</sub>	Μ̈́S												
14C	LG												
Bact	CPA							¥	Y	¥	⊁	*	<b>}</b>
Z00	CS					7						⊁	
DMS	RS												>-
ומ	AT							Υ	Ϋ́	Y	Υ	<b>&gt;</b>	>
TCO2	Qſ					λ.			Y	Y	Y		<b>&gt;</b> -
Hydr	WB												
Halo	JB												
OŽN	TF												
$SF_6$	FC					Ϋ́	Y	Ϋ́	¥	¥	⊁	Y	λ
Sai	JR	Υ	Y	Y	Y	Å	Ϋ́	Y	Ϋ́			λ	
Nuts	ŢĴ					λ	Y	Ϋ́	×	Y	>	<b>&gt;</b>	<b>&gt;</b>
AFC Phyto	SA					Y			¥	¥	<b>&gt;</b>	<b>*</b>	Y
Phyto	MHL					155			154	153	152	151	150
HPLC	MHL					<b>×</b>			Ϋ́	Ϋ́	>	¥	<b>}</b>
Chi	MHL					٠	X	Ϋ́	Ϋ́	Y	Y	λ	>
Bottle		4R	8R	12R	20R	2T	6T	10T	14T	16T	18T	22T	24T
Depth		200	400	300	250	200	150	001	20	40	25	10	G

CTD41-2/7/98, 17:15-17:30, 59°59'56"N, 20°04'02"W, Feature to N, 3 bottles fired at 13 m

CTD42-2/7/98, 17:50-19:40, 59°59'56"N, 20°04'02"W, DEEP #1

43		$\exists$	7	1		Ţ		_			[	*****	
DMSe	PL												
<u>Z</u>	GM	<b>&gt;</b> -		٨									
ပ္	TG	*											
Bact	CPA	¥	Y	ž	>	>	*	⋆	Ϋ́				
Z00	S												
DMS	RS	***************************************									-		
Ω	AT	Y	¥	>-	>-	<b>}</b>	>						
$TCO_2$	Ωſ		Y	<b></b>		Y	Y		Y				
Hydr	WB	Ϋ́	Ϋ́	λ		Y		Y					>
Halo	JB	Ϋ́	Y	Ϋ́	<b>*</b>	λ	Y						>
N <sub>2</sub> O	TF	Y	>	¥	¥	٠	λ	¥					>
$SF_6$	FC	Ÿ	<b>,</b>	Y	Ϋ́	Y	Ÿ	Ÿ					^
Sal	JR		<b>~</b>			¥	Ϋ́	Y	Y	Ă	Å	Å	Λ
Nuts	Ľ	Y	λ	Ž	Ϋ́	Y	Ϋ́	¥	Y	Y	Å	Å	Λ
AFC Phyto	SA	Ý	Ÿ	Ÿ	Ϋ́	Ϋ́							
Phyto	MHL	173	174	175	176	177	178	179					
HPLC	MHL	×	>	¥	<b>~</b>	<b>&gt;</b>	>	Ϋ́					
Chl	MHL	Y	>-	<b>*</b>	<b>-</b>	٨	Ϋ́	Ϋ́					<u> </u>
Bottle		24T	22T	18T	16T	14T	10T	T9	20R	12R	8R	4R	3.7
Depth		0	10	20	30	50	100	200	500	Г	Г		

CTD43 - 2/7/98, 21:26-23:57, 59°53'04"N, 20°21'48"W, Outside, DEEP #2

Lyase	37.1	CIVI														>
Chi	760	ĭ									Y	>-	>	ĭ	Υ	>
DMSe	ă	T.C.														
Z <sub>S</sub> 1	y (2)	74									X	<b>&gt;</b>	^	Υ,	Y	<b>~</b>
14C	1.0	3														
Bact	VDV	5					Y		>	- }	Y	>-	>	Ţ		>-
Z00	٧	3										>-	>	1	X	<b>≻</b>
DMS	PS															
Ω	AT									>	1		٨	٠ >	I	>-
TCO <sub>2</sub>	1								***************************************							
Hydr	WB		-						-							Y
Halo	E,								-				>-			Y
N <sub>2</sub> O	TF	>	į						>	^	*		>-		3	Y
$SF_6$	55	>						>-	>	>	\ \ \ \	λ	X	>	;	Y
Sal	JR.	>	\  >	,   }	· >	·   >	-	>-	X	>				<b>&gt;</b>		
Nuts	Ţĵ												×		,	λ
AFC Phyto	SA										1	¥	<b>&gt;</b>	٨		X
Phyto	MHL												181		100	180
HPLC	MHL												¥		>	-
Chl	MHL												Y		>	I
Bottle		2	48	8R	12R	20R		0	01	14	1,4		81	22	2.4	7.4
Depth		2760	2000	1500	1000	500	000	007	100	50	30	3	70	10	-	2

CTD44 - 3/7/98, 01:20-03:06, 59°46'22"N, 20°39'11"W, Edge, DEEP #3

_	_	-	_		••••	_		_		•	_	_	_			_		
Lyase	0),	CIVI	<u>&gt;</u>	>	, ,	1	⊁	>	-									
Fe	Ę	f.		-								;	X	>	, ,	X	>	
DMSe	2	7.							-									
N <sub>S1</sub>	280	5	>-		>	4												
<sup>14</sup> C	2	3										,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,						
Bact	۷۵۷	5 3	<b>&gt;</b>	>	>	1 ;	Y	>	, ,	¥		>	I					
Z00	20	3	<b>&gt;</b> -	>	<b> </b> >	, ;	X	>		ĭ				-				
DMS	28																	
ă	AT		×	٨	>	, ,	Y	7	>	ľ			-					
TCO2	1								>	1		>	¥					
Hydr	WB	) }	Y	~	>			<b>&gt;</b>	>	*				******				Α.
Halo	B			>-	>			<b>≻</b>					-					
N <sub>2</sub> O	TF	>	I	>-	Y	>	-	>-	>	Ý	>-							Ϋ́
${ m SF}_6$	<u> </u>	>	<u> </u>	>	Ă	>	1	>	٨	Ţ	×							>
Sail	JR			Y				>-	>	•	>	>	**	Į,	Y		Y	>-
Nuts	ŢĴ	>	1	٨	>-	>	1	<b>\</b>	<b>*</b>		¥	>	1.7	ľ	Y	1.7	ĭ	Y
AFC Phyto	SA																	
Phyto	MHL	182	707	183	184	185		186										
HPLC	MHL	>		Y	>	<b>~</b>		Y	>	;	Y							
5	MHL	٨	,	λ	Υ	<u>,</u>		X	>		Υ							
Bottle		24		777	18	16		14	10		٥	20R	120	177	SR.	45	4	7
Deptin		C	١	97	20	30		25	100	900	700	500	1000	2000	1500	2000	2000	2790
						-		_	-						_		_	_

CTD45 -- 3/7/98, 04:40-06:34, 59°39'41"N, 20°56'58"W, Outside, DEEP #4

DMSe	ΡΓ												
<u>z</u>	GM												
ئ ئ	97	٨											
Bact	CPA	>	⊁	X	Y	Y	χ.		X				
Z00	CS												
DMS	RS												
Ω	AT												
TCO2	Ωſ												
Hydr	WB												
Halo	JB												
O <sup>2</sup> N	TF												
$\mathrm{SF}_6$	FC	Y	Υ	γ	Ă	Y	Υ	Y					<b>&gt;</b>
Sal	JR		Y			<b>&gt;</b>	¥	<b>&gt;</b>	Y	Υ	Y	Y	>
Nuts	TJ												
AFC Phydo	SA												
Phyto	MHL	187		188									
HPLC	MHL	Ϋ́		Y									
Chl	MHL	<b>}</b>		Υ									
Depth Bottle		24	22	18	16	14	10	9	20R	12R	8R	4R	,
Depth		0	10	20	30	20	100	200	200	1000	1500	2000	2804

CTD46 - 3/7/98, 08:14-10:14, 59°32'42"N, 21°13'48"W, Outside, DEEP #5

1			Т	<del></del>	1	_		_	_	_	1	1	_
Lyase	MS	Y											
DMSe Lyase	PL												
N <sub>S1</sub>	ВM	>		>							_		
Jы C	LG	>		>	≻								
Bact	CPA	>-	>	¥	Ϋ́	Ϋ́	¥		>				
Z00	CS	Y	7	¥	Y	Y	¥						
DMS	RS												
Ĭ	AT	Y	¥	>-	٨	¥	≻						
TCO2	JD												
Hydr	WB												
Halo	JB	Å	Υ	Ă.		Ă				-			
O <sub>Z</sub> N	TF	Å	¥	Y	Ă	Y	¥	Ý					Ϋ́
$\mathrm{SF}_6$	FC	¥	Y	λ	λ	¥	×	Ă					<b>&gt;</b> -
Sal	JR		Å			>	>-	۲	Ϋ́	Ϋ́	Ă	Ϋ́	Å
Nuts	ŢJ	Y	Y	7	≻	Υ	7	>-	Ϋ́	Ă	Y	Ϋ́	Ă
AFC Phyto	SA	<b>&gt;</b>	¥	Υ	Y	Ý	>	Ϋ́					
Phyto	MHL	189	190	161	192	193							
HPLC	MHIL	٠	Ϋ́	Ϋ́	¥	Ϋ́	¥	٨					
Chi	MHIL	<b>&gt;</b>	Ϋ́	Ý	Ϋ́	>	>	>-					
Bottle		24	22	18	16	14	0	9	20R	12R	8R	4R	2
Depth		0	10	20	30	50	100	200	200	1000	1500	2000	2840

CTD47-3/7/98, 11:42-13:37, 59°25'43"N, 21°31'24"W, Salty Feature, DEEP #6

DMSe	Ы	1					T						
<sup>15</sup> N DMSe	NU NU	†					-			+	-		
) <sub>1</sub> C	٢											-	
Bact	CPA	T	· >	· >	Ý	\ \ \	- 						
Z00	S												
DMS	RS												
á	AT												
TCO2	Q.		Y		<b>~</b>	>			Y				
Hydr	WB												
Halo	JB	Y		<u>۲</u>									
OzN	TF												
$\mathrm{SF}_6$	FC	X	Ϋ́	Y	Ϋ́	7	¥	>-					¥
Sal	JR		Y			X	<b>&gt;</b>	}	X	<b>}</b>	γ	\- \-	Y
Nuts	IJ	>		Y									
AFC Phyto	SA												
Phyto	MHL	194		195									
STAH	THM	λ		Ϋ́									
Chi	THM	Ä		Ϋ́									
Bottle		24T	22T	18T	191	14T	101	6T	20R	12R	8R	4R	2T
Depth		0	10	20	30	20	100	200	500	1000	1500	2000	2850

CTD48-3/7/98, 14:53-16:46, 59°18'45"N, 21°48'21"W, Outside, DEEP #7

		<del></del>	<del></del>	<del></del>	т-	<del>,</del>	_	·	,	_	<del>,</del>	<del></del>	_	
DMSe		ы	2											
N <sub>S</sub> 1		Š												
14C	<b>)</b>	2												
Bact		CPA	>	, >	γ	<u>\</u>	· >	À						
Zoo		S												
DMS		RS												
Q		AT	>	>	<b>&gt;</b>	>	>	<b> </b> >-	***************************************					
TCO,	1	2			¥		>-	>-		-				
Hydr	,	WB	¥	>~	λ	<b>&gt;</b>	Y							>
Halo		E G	×	<u></u>	×	Ϋ́	Ϋ́							
O'N		TF	7	¥	λ	Y	Y	Υ	Å					Y
$SF_6$		FC	>	\	λ	Ϋ́	Å	Ϋ́	¥					Ÿ
Sal		Æ		<b>~</b>			¥	⋋	X	¥	¥	λ	Ÿ	Υ
Nuts		£	≻	<u>}</u>	۲,	¥	Υ	Ϋ́	7	>-	Y	Υ	Ϋ́	Y
AFC	Phyto	SA	Ă	¥	X	Y	Ķ							
Phyto		MHL	196	197	198	199	200							
HPLC		THM	Å	<b>*</b>	Ϋ́	Y	Å	Ϋ́	Ϋ́					
Ch1		MHE	· Х	¥	Ϋ́	Ă	Ϋ́	Å	Ă			·		
Bottle			24T	22T	181	16T	14T	10T	ET.	20R	12R	8R	4R	2T
Depth			0	10	20	30	50	100	200	200	1000	1500	2000	2800

CTD49 3/7/98, 18:22-20:15, 59°11'38"N, 22°05'20"W, Outside, DEEP #8

0)													
DMS	PL												
<sup>14</sup> C 15N DMSe	ВМ												
¢C	T.G												
Bact	CPA	Å	Ă	Å	Y	Ă	Ā						
Z00	CS												
DMS	RS												
ΙΟ	AT												
TCO2	JD												
L	WB												
Halo	JB	Ā		٠									
N <sub>2</sub> O	TF	Å		Å		Y	Å						Å
SF6	FC												
Sal	JR		Ϋ́			Y	Y	Y	λ	Ā	Å	Y	Y
Nuts	TJ	Ϋ́		Ϋ́									
AFC Phyto	SA												
Phyto	MHL	201		202									
НРЬС	MHL	Ϋ́		λ									
СЫ	THM	Y		Å									
Bottle		24T	22T	18T	16T	14T	10T	ET	20R	12R	8R	4R	2T
Depth		0	10	20	30	20				0001	1500	2000	2783

CTD50 - 3/7/98, 22:20-00:07, 59°04'31"N, 22°21'53"W, Outside, DEEP #9

							П	_					$\neg$
Lyase	MS	λ	Y	Y	Ÿ	Ϋ́							
DMSe	PL	۱ ۲	Y	λ	λ	Y	~						Y
Z <sub>SI</sub>	GM												
Chi	1.G	Y		λ	Ý								
Bact	CPA	Y	Y	Å	Ÿ	Ϋ́	Y		Ϋ́				
Z00	CS	Υ	λ	Ϋ́	λ	Ϋ́	Y						
DMS	RS												
ם	AT	Ϋ́	Å	Ý	Ă	Å	Ϋ́						¥
TCO2	JD												
Hydr	WB	҂											
Halo	JB												
O <sub>2</sub> N	TF	¥	Y	Y	Ā	λ	Ÿ	Y					Ÿ
SF <sub>6</sub>	F.		Ÿ										
Sal	JR		٨			Ϋ́	Ϋ́	Y	Ϋ́	Ā	Ϋ́	Y	Y
Nuts	Ω	Ϋ́	Ϋ́	Ă	¥	>-	¥	Y	Ă	Ϋ́	Ă	٠.	λ
AFC Phyto	SA	<b>&gt;</b>	¥	×	>-	>-	Ÿ						
Phyto	MHL	203	204	202	206	207							
HPLC	MHL	Y	Y	λ	λ	Y	Ý	Å					
Chi	MHL	Ϋ́	Ϋ́	Ÿ	Y	Y	¥	¥					
Bottle		24	22	18	91	14	10	9	20R	12R	8R	4R	2
Depth		0	10	20	30	50	100	200	500	1000	1500	2000	2770

CTD51 - 4/7/98, 04:42-04:52, 59°20'22"N, 21°44'43"W, Outside

·			_			_	 		
DMSe	PL								
N <sub>St</sub>	ВМ								
14C	91								
Bact	CPA								
Zoo	S	Ϋ́	Ϋ́						
DMS	RS								
Q	AT								
TCO2	JD								
Hydr TCO2	WB			Y					
Halo	JB								
N <sub>2</sub> O	TF								
$SF_6$	PN								
Sat	JR								
Nuts	TJ								
AFC Phyto	SA								
	MHL								
HPLC Phyto	MHI						-		
Chl	MHL						_		
Depth Bottle		2	4R	9					
Depth		4	4	4					

CTD52 - 4/7/98, 05:15-05:34, 59°20'22"N, 21°44'43"W, Outside

<b></b>	<del></del>	,								······			
DMSe	PL												
N <sub>S1</sub>	GM	Ÿ		Y									٨
Chl	TG		Ϋ́	Y		Y	¥		¥				Ϋ́
Bact	CPA												
Zoo	cs									Ϋ́	Å		
DMS	RS												
Ω	AT												
$TCO_2$	Ωſ												
Hydr	WB												
Halo	JB	Y	X		Ϋ́	Ÿ		¥	Ϋ́			҂	
N <sub>2</sub> O	TF												
$\mathrm{SF}_6$	PN												
Sal	JR												
Nuts	IJ	Å	Ϋ́	¥		Ă		Ă				Ă	
AFC Phyto	SA	Y	Ϋ́	Ÿ			Ÿ		Ÿ	Y	Y	Ϋ́	
Phyto	MHL												
HPLC	MHL												
СЫ	MHL												
Bottle		2	9	8R	10	14	12R	16	24	20R	4R	18	22
Depth		50	30	20	20	17	11	11	4	4	4	0	0

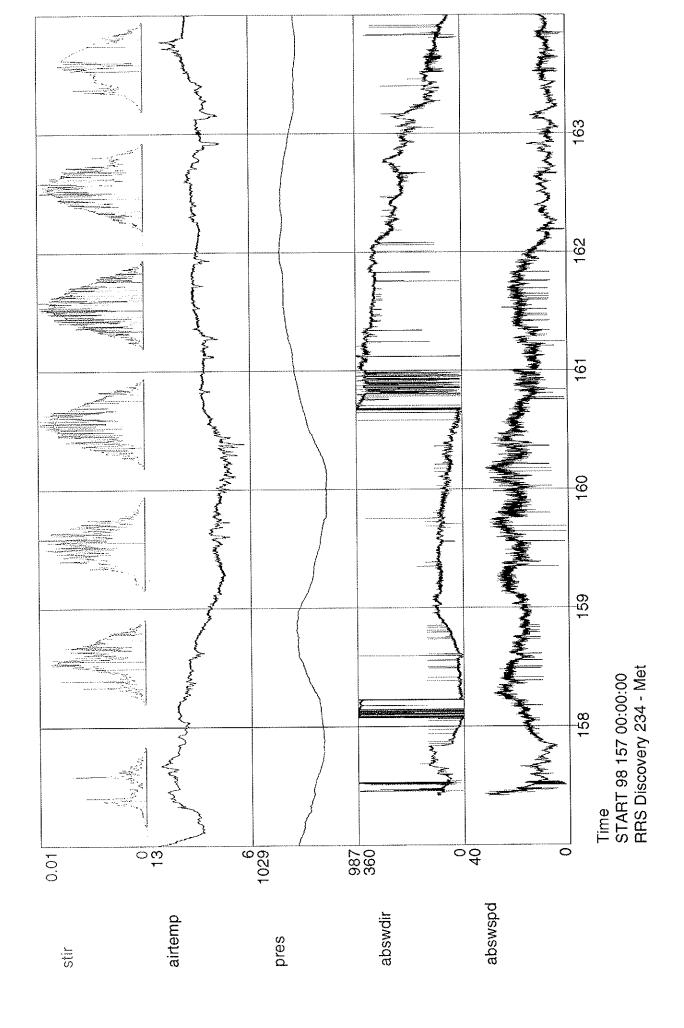
CTD53 - 4/7/98, 18:34-18:42, 59°20'30"N, 21°55'28"W, Outside

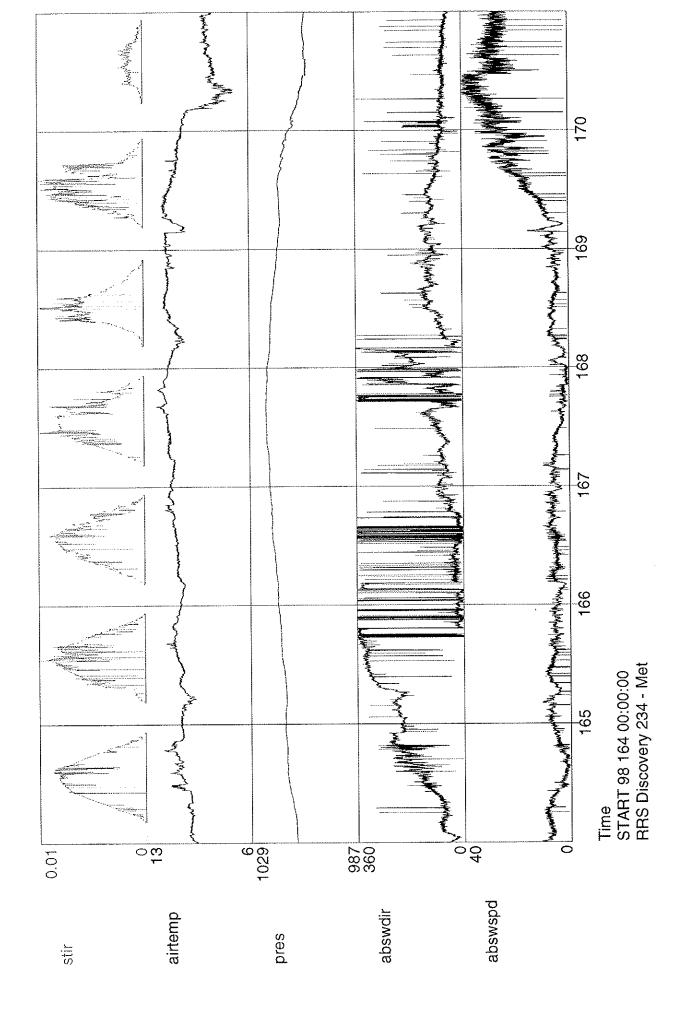
		_		 <del></del>		1	7	_	 	 	1
Lyase	MS	;	X.								
DMSe	PL										
<sup>14</sup> C   <sup>15</sup> N DMSe Lyase	GM										
14C	57										
Bact	CPA										
Zoo	CS	<b>&gt;</b>									
DMS	RS										
G	AT										
Hydr TCO2	Ωſ	-				].					
Hydr	WB										
Halo	JB										
O <sub>2</sub> N	TF										
SF <sub>6</sub>	FC										
Sal	JR								 		
Nuts	Tì							_			***************************************
AFC	SA		Y							_	
Phyto	MHL		208								-
HPLC	MHL		Y								
Chi	MHL		Ϋ́								
Depth Bottle		18T	22T								
Depth		0	0								***

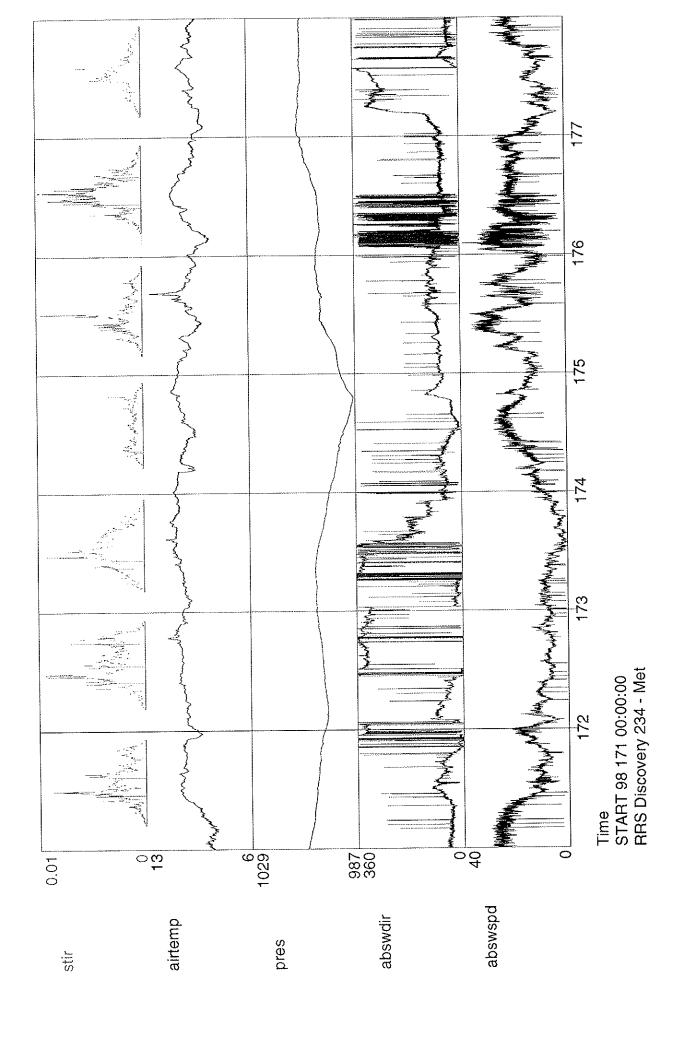
CTD54 - 6/7/98, 06:12-08:35, 54°24'30"N, 14°50'10"W, Outside

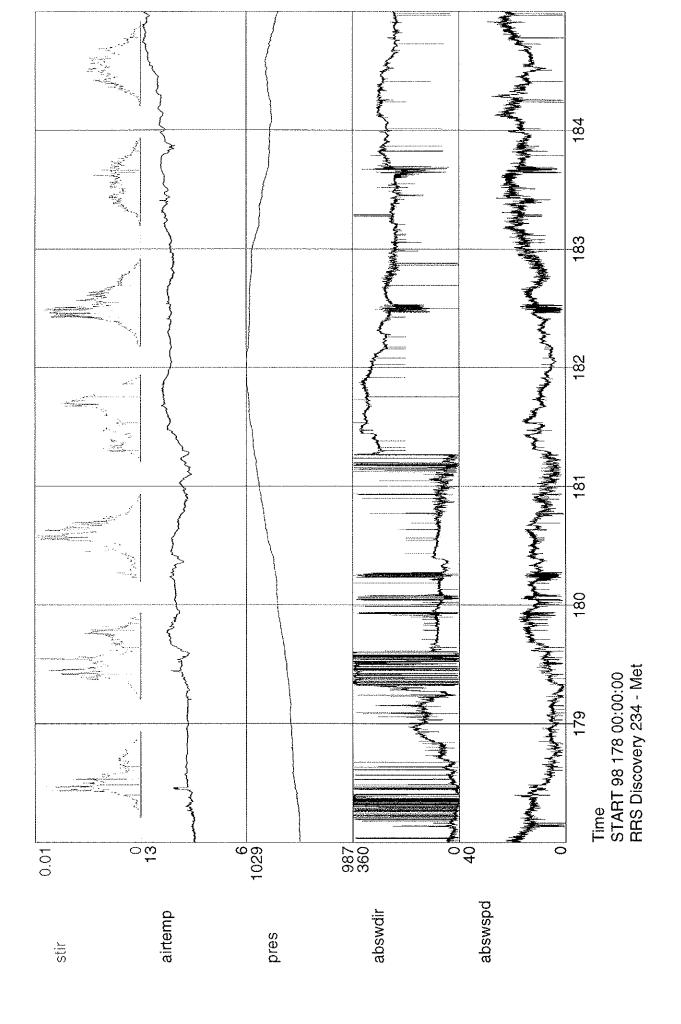
DMSe	į	P.L.												
$z_{\rm si}$		C.W												
14C		FG												
Bact		CPA												
Zoo	ļ	S												_
DMS		RS												
Q _		ΑŢ												
TCO2		ŭ												
Hydr		WB												
Halo		JB											Υ	
N2O		TF	λ.		Y		҂							
SF6		FC	Ϋ́		>-		<b>&gt;</b>		٨	٨	>		Y	<b>&gt;</b>
Sal		R	Y	Y	>	٨	Y	Y	Y	¥	¥	Х	≻	>
Nuts		TJ	¥	Y	Y	Y	¥	⊁	λ	Y	Υ	Ϋ́	Y	>-
AFC	Phyto	SA												
Phyto		MHL												
HPLC		MHL												
Chl		MHL												
Bottle	******		2	4R	9	8R	10	12R	14	16	18	20R	22	24
Depth	!		2710	2710	2710	2710	2710	2710	2710	2710	2710	2710	2710	2710

### METEOROLOGICAL DATA - WEEKLY PLOTS

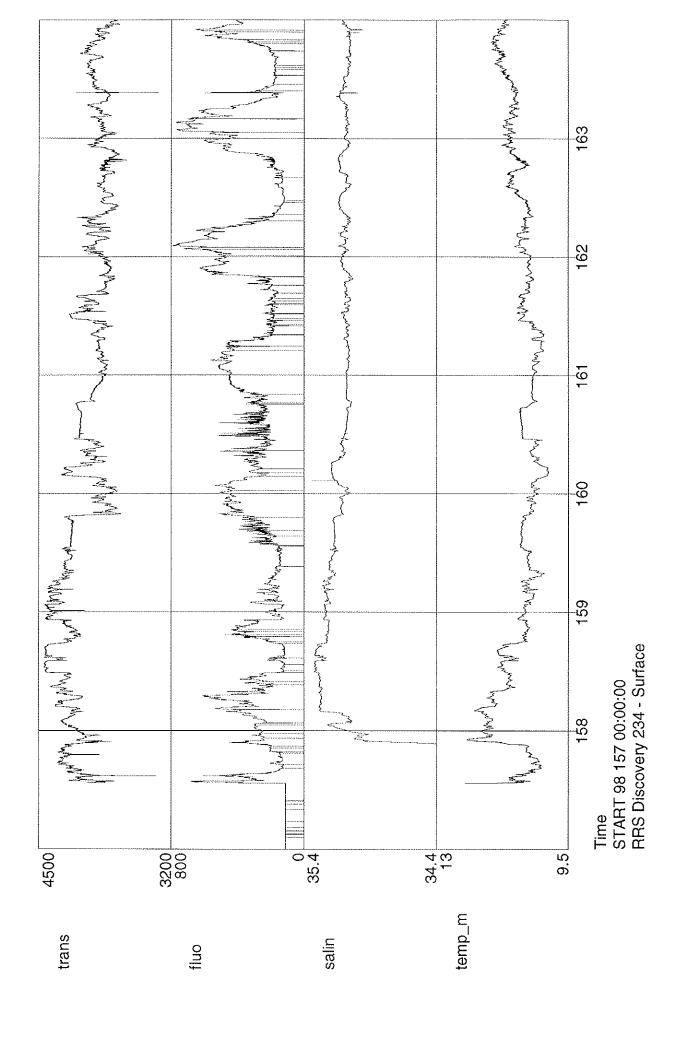


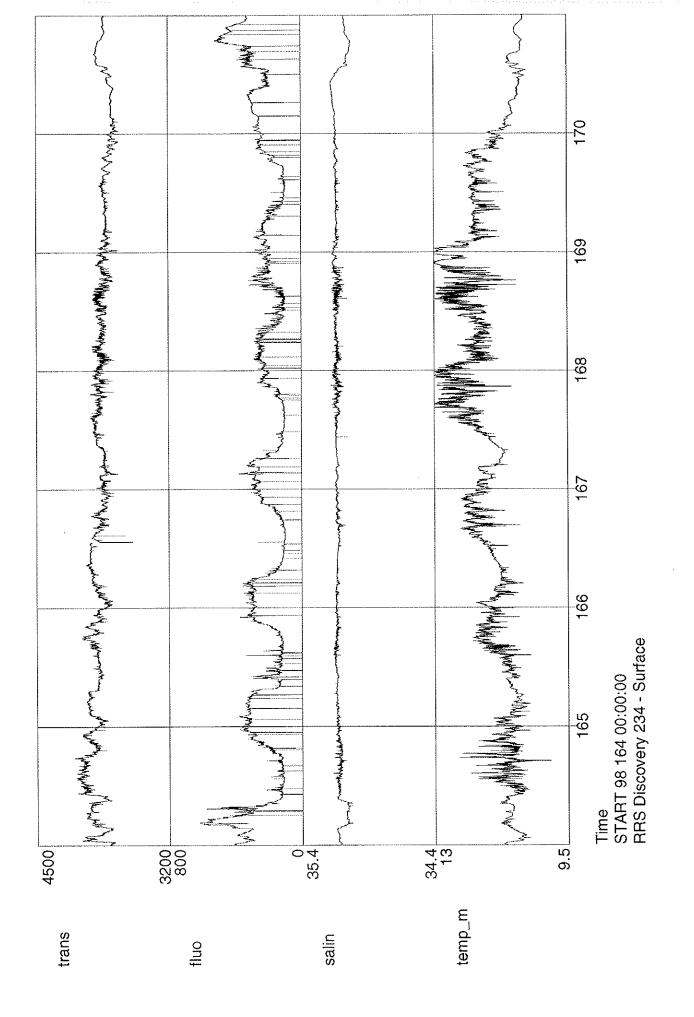


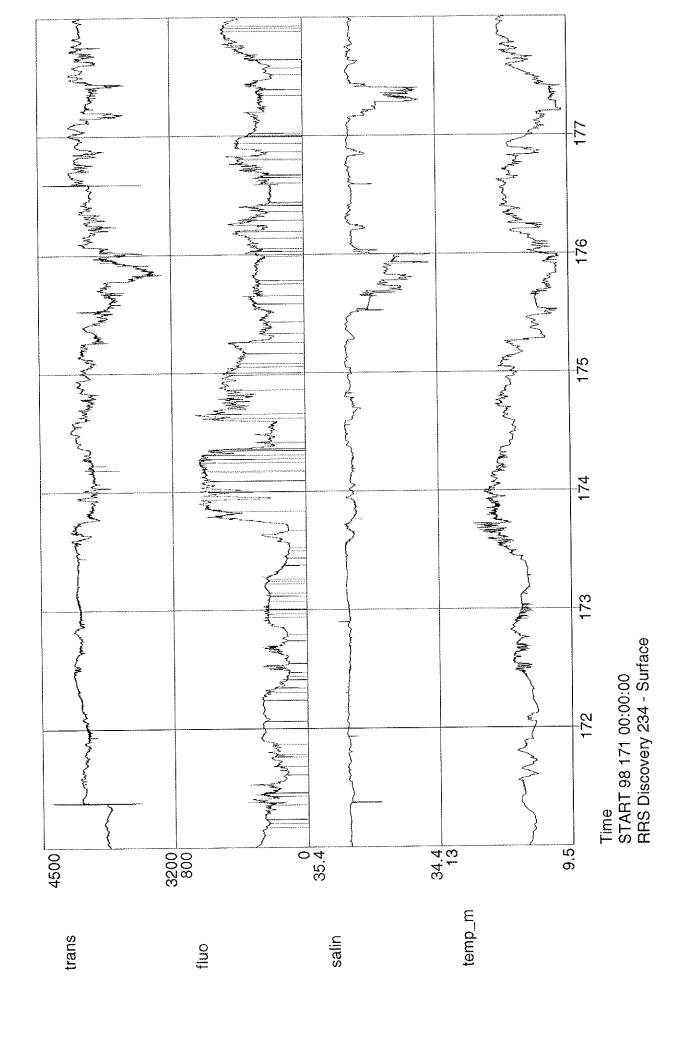


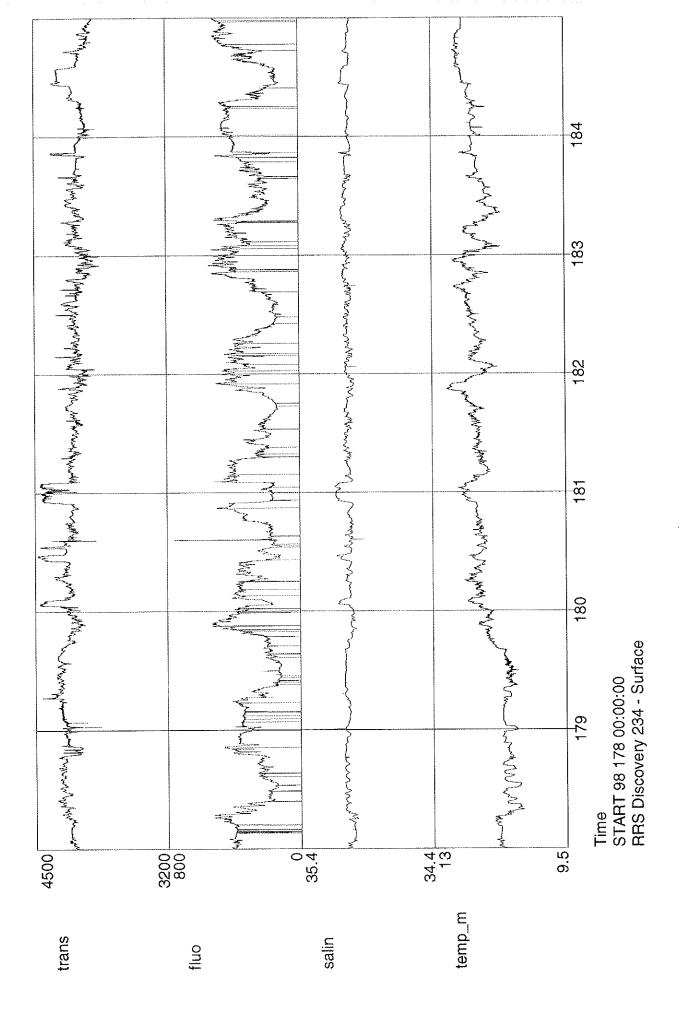


# WEEKLY UNDERWAY PLOTS - TEMPERATURE, SALINITY, TRANSMISSOMETER AND FLUORESCENCE









## **CTD PLOTS**

