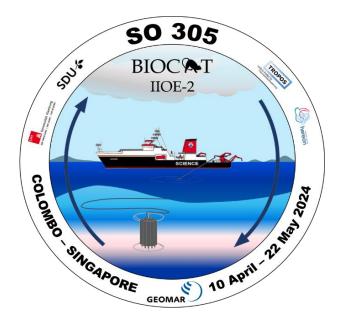
SONNE-Berichte

Biogeochemistry/atmosphere processes in the Bay of Bengal: A contribution to the 2nd International Indian Ocean Expedition

Cruise No. SO305

10 April – 22 May 2024 Colombo (Sri Lanka) – Singapore (Singapore) BIOCAT-IIOE2



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1 Cruise Summary

1.1 Summary in English

The overarching goal of SO305 BIOCAT-IIOE2 was to quantify key (micro)biological processes in the water column and ocean/atmosphere exchange fluxes to assess their impacts on the oxygen minimum zone (OMZ) of the Bay of Bengal. To this end, we conducted a measurement campaign with the research vessel (RV) SONNE from 10 April to 22 May 2024 (SO305) as part of the BIOCAT-IIOE2 project, covering the main carbon and nitrogen cycle processes and physical processes in the water column. The oceanic measurements were complemented by an intensive atmospheric measurement program to investigate the effects of atmospheric inputs on water column processes. During SO305 the GEOMAR Helmholtz Centre for Ocean Research Kiel, the University of Hamburg, the Helmholtz Centre Hereon (Geesthacht), the Leibniz Institute for Tropospheric Research (TROPOS, Leipzig), the University of Oldenburg and the University of Southern Denmark (SDU, Odense, DK) were collaborating. A team of 39 scientists, students and technicians made measurements in the water column and in the atmosphere at 33 regular CTD stations and five 24h-stations along a cruise track from the eastern equatorial Indian Ocean to the central Bay of Bengal. The results of SO305 BIOCAT-IIOE2 will contribute to a significantly improved assessment of the future impacts of global climate change and pollution for the ecosystems and the OMZ of the Bay of Bengal.

1.2 Zusammenfassung

Das übergeordnete Ziel von SO305 BIOCAT-IIOE2 war, die wichtigsten (mikro)biologischen Prozesse in der Wassersäule und die Austauschflüsse zwischen Ozean und Atmosphäre zu quantifizieren, um ihre Auswirkungen auf die Sauerstoffminimumzone (SMZ) des Golfs von Bengalen zu bewerten. Zu diesem Zweck haben wir vom 10. April bis zum 22. Mai 2024 eine Messkampagne mit dem Forschungsschiff (FS) SONNE durchgeführt (SO305), die die wichtigsten Prozesse des Kohlenstoff- und Stickstoffkreislaufs sowie physikalische Prozesse in der Wassersäule erfasste. Die ozeanischen Messungen wurden durch ein intensives atmosphärisches Messprogramm ergänzt, um die Auswirkungen der atmosphärischen Einflüsse auf die Prozesse in der Wassersäule zu untersuchen. Während SO305 haben das GEOMAR Helmholtz-Zentrum für Ozeanforschung Kiel, die Universität Hamburg, das Helmholtz-Zentrum Hereon (Geesthacht), das Leibniz-Institut für Troposphärenforschung (TROPOS, Leipzig), der Universität Oldenburg und der University of Southern Denmark (SDU, Odense, DK) zusammengearbeitet. Ein Team von 39 Wissenschaftlern/innen, Studenten/innen und Technikern/innen führte Messungen in der Wassersäule und in der Atmosphäre an 33 CTD-Stationen und fünf 24-Stunden-Stationen entlang einer Fahrtroute vom östlichen äquatorialen Indischen Ozean zum zentralen Golf von Bengalen durch. Die Ergebnisse von SO305 BIOCAT-IIOE2 werden dazu beitragen, die künftigen Auswirkungen des globalen Klimawandels und der Umweltverschmutzung auf die Ökosysteme und die SMZ des Golfs von Bengalen wesentlich besser einschätzen zu können.

2 Participants

2.1 Principal Investigators

Name	Institution
Bange, Hermann, Prof. Dr.	GEOMAR
Gaye, Birgit, Dr.	Univ. Hamburg
Löscher, Carolin, Prof. Dr.	SDU
van Pinxteren, Manuela, Dr.	TROPOS



Fig. 2.1 Participants of SO305 BIOCAT-IIOE2 on 09 April 2024; in front of RV SONNE before departure from Colombo, Sri Lanka. In the front, sitting, from left to right: Qingwen Zhong, Jon Roa, Gesa Schulz, Anja Conventz, Isabell Schlangen, René Rabe, Arun Babu Suja, Shravan Deshmukh, Lubrina Nielsen, Jule Ploschke and Riel Ingeniero. In the back, standing, from left to right: Hermann Bange, Felix Duerkop, Marcel Sommer, Antoine Barbot, Albert Firus, Ina Stoltenberg, Victor Fernández-Juárez, Mario Müller, Isabell Hentschel, Birgit Quack, Sandra Golde, Theresa Barthelmeß, Paula Eisnecker, Clara McKellar, Martha Gledhill, Laurenz van Bonn, Mats Jacobsen, Leon Schmidt, Hendrik Großelindemann, Jannis Usinger, Julia Mickenbecker, Hendrik Feil, Kira Lange, Dennis Booge, Sönke Rolfes, Tjark Andersch, Rena Czeschel and Leandro Nazzari.

Name	Discipline	Institution
Bange, Hermann, Prof. Dr.	Trace gases / Chief Scientist	GEOMAR
Schmidt, Leon	Nutrients, O ₂	Hereon
Usinger, Jannis	Nutrients, O ₂	GEOMAR
Lange, Kira	Nutrients, O ₂	GEOMAR
Czeschel, Rena, Dr.	CTD, microstructure, Argo float	GEOMAR
	deployment, mooring deployment	
Müller, Mario	CTD, microstructure, Argo float	GEOMAR
	deployment, mooring deployment	
Zhong, Qingwen	CTD, microstructure, Argo float	GEOMAR
	deployment, mooring deployment	
Großelindemann, Hendrik	CTD, microstructure, Argo float	GEOMAR
	deployment, mooring deployment	
Duerkop, Felix	CTD, microstructure, Argo float	GEOMAR
	deployment, mooring deployment	CEOMAD
McKellar, Clara	CTD, microstructure, Argo float	GEOMAR
Stoltenberg, Ina, Dr.	deployment, mooring deployment N ₂ O, CH ₄ , NO	GEOMAR
Van Bonn, Laurenz	N ₂ O, CH ₄ , NO	GEOMAR
Hentschel, Isabell	N_2O, CH_4 N ₂ O, CH ₄	GEOMAR
Eisnecker, Paula		GEOMAR
Sommer, Marcel	CO, N ₂ O, CH ₄ , N ₂ O/CO underway CO, N ₂ O, CH ₄ , N ₂ O/CO underway	GEOMAR
Ingeniero, Riel	NO	GEOMAR
Barthelmeß, Theresa, Dr.		GEOMAR
Golde, Sandra	POM, DOC, drifting sediment traps POM, DOC, drifting sediment traps	GEOMAR
Roa, Jon	POM, DOC, drifting sediment traps	GEOMAR
Barbot, Antoine		GEOMAR
Gledhill, Martha, Dr.	POM, DOC, drifting sediment traps Trace elements/metals	GEOMAR
Firus, Albert	Trace elements/metals	GEOMAR
	Trace elements/metals	GEOMAR
Conventz, Anja Quack, Birgit, Dr.	Halocarbons	GEOMAR
Ploschke, Jule	Halocarbons	GEOMAR
	Halocarbons	
Mickenbecker, Julia		GEOMAR GEOMAR
Booge, Dennis, Dr. Feil Hendrik	DMS, isoprene	GEOMAR
Feil, Hendrik	DMS, isoprene	
Rolfes, Sönke	DMS, isoprene, meta genomics	U. Oldenbu
Schulz, Gesa, Dr.	¹⁵ N isotopes, suspended mater	U. Hamburg
Andersch, Tjark	¹⁵ N isotopes, suspended matter	U. Hamburg
Nazzari, Leandro	¹⁵ N isotopes, suspended matter	U. Hamburg
Schlangen, Isabell	Microbial processes, meta genomics	SDU
Jacobsen, Mats	Microbial processes, meta genomics	SDU

2.2 Scientific Party

Nielsen, Lubrina	Microbial processes, meta genomics	SDU
Fernández Juárez, Victor, Dr.	Microbial processes, meta genomics	SDU
Rabe, René	Aerosols, atm. trace gases	TROPOS
Babu Suja, Arun, Dr.	Aerosols, atm. trace gases	TROPOS
Deshmukh, Shravan	Aerosols, atm. trace gases	TROPOS

2.3 Participating Institutions

Helmholtz-Zentrum für Ozeanforschung Kiel
Helmholtz-Zentrum Geesthacht
University of Southern Denmark, Odense, Denmark
Leibniz-Institut für Troposphärenforschung, Leipzig
Universität Hamburg
Universität Oldenburg

3 Research Program

3.1 Description of the Work Area

3.1.1 Bay of Bengal

The Bay of Bengal (BoB, located between 05°-23°N and 80°-95°E) forms the north-eastern basin of the Indian Ocean. The BoB is characterized by a unique environmental setting which shows a remarkable seasonality driven by the Asian monsoon system with a summer monsoon season (SW monsoon) from June-August and a winter monsoon season (NE monsoon) from December-February. The major physical drivers of the environmental setting of the BoB are summarized in Fig. 3.1. The Asian monsoon system leads to

- (i) pronounced rainfall and river discharge with maximum freshwater inputs in September by the Ganges/Brahmaputra and Irrawaddy river systems in the northern BoB,
- (ii) a seasonally reversing upper ocean circulation and
- (iii) a severe atmospheric pollution (known as the 'South Asian Brown Cloud') which lasts over the BoB from November to May.

Moreover, the BoB hosts one of the world-wide most intense oxygen minimum zones (OMZ) with persistent minimum oxygen concentrations as low as $0.01 \mu M$ (Bristow et al., 2017).

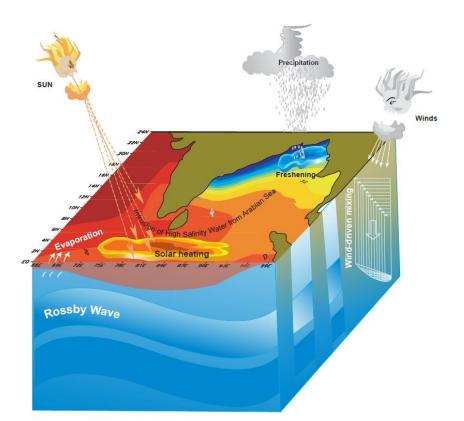


Fig. 3.1 Schematic representation of the local and remote physical forcings that influence the water column of the BoB. Colour shading depicts the climatological monthly mean salinity for August. Local forcing that affect the BoB are precipitation, river runoff, solar heating and mixing by monsoon winds. The remote forcing are the intrusions of high salinity waters from the Arabian Sea and propagation of Rossby waves. (Fig. is taken from Narvekar and Prasanna Kumar, 2014).

3.1.2 Eastern Equatorial Indian Ocean

The eastern part of the equatorial Indian Ocean is located at the southern boundary of the BoB between 05°N-10°S and 80-95°E and separates the BoB from the south-eastern Indian Ocean. The equatorial circulation of the Indian Ocean is characterized by annual mean winds that are westerly along the equator rather than easterly. These winds favour downwelling at the equator rather than upwelling as observed in the equatorial Pacific and Atlantic Oceans (Phillips et al., 2024). Moreover, seasonal reversals of the winds along the equator lead to strong semi-annual variability, with strong eastward currents, known as Wyrtki jets, dominating the upper 100 m during the transitions between the NE and SW monsoons (Phillips et al., 2024). There is no permanent equatorial undercurrent, i.e., no distinctive core of eastward flow in the thermocline. An undercurrent-like structure does appear in February-March near the end of the NE monsoon following a two-month period of sustained easterly winds along the equator, but this feature is transient and quickly disappears with the onset of westerly winds during the following monsoon transition season. A weaker and more variable undercurrent appears in August–September (Phillips et al., 2014).

3.2 Aims of the Cruise

The overarching goal of SO305 BIOCAT-IIOE2 was to quantify the key (micro)biological processes in the water column and ocean/atmosphere exchange fluxes in order to assess their impacts on the OMZ of the Bay of Bengal. The specific goals of the cruise were:

- To decipher the physical and biogeochemical setting of the water column,
- To identify the physical and biogeochemical/microbial processes which are crucial for the, development and maintenance of the OMZ,
- To assess the efficiency of the biological pump,
- To quantify the fluxes of climate-relevant trace gases across the ocean/atmosphere interface,
- To estimate the fluxes of trace metals and nutrients to the ocean from the ocean boundaries (i.e. atmosphere, continent and sediments) and
- To characterize the species and amount of natural and anthropogenic constituents (incl. trace gases and aerosols) in the marine boundary layer of the Asian outflow and their effects on the self-cleaning capacity (i.e. oxidizing efficiency) of the atmosphere.

3.3 Agenda of the Cruise

The regular, repetitive work at 38 stations included the use of the CTD/Rosette (CTD/Ro, from the water surface to the sea floor at a maximum water depth of 4500 m), microstructure measurements (with a free-falling microstructure probe up to 200 m water depth) and the use of GoFlo water samplers (up to a water depth of 500 m). At five 24h-stations, the regular station program was supplemented by zodiac deployments to sample the uppermost meter of the water column, the use of a submersible pump (up to a water depth of 150 m) and the deployment of drifting sediment traps (which were picked up again after 48 hours). In addition to the station work, continuous measurements were carried out in the atmosphere (trace gases, aerosols) and in the surface water (dissolved trace gases and sampling for trace metals with a Towed-fish). Moreover, we deployed a long-term deep-sea mooring at the equator and deployed two Argo floats in the southern and central BoB. The cruise track of SO305 BIOCAT-IIOE2 is shown in Fig. 3.2.

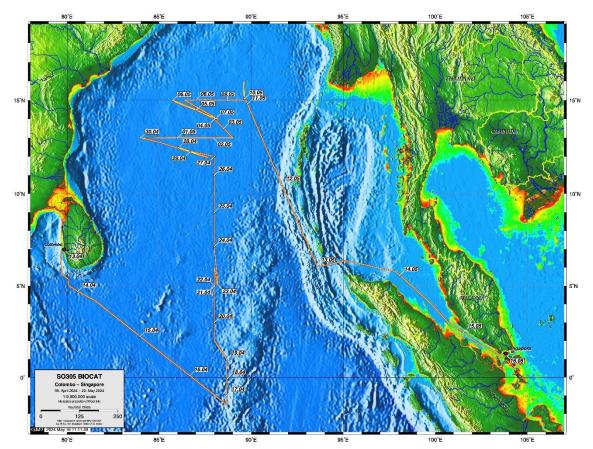


Fig. 3.2 Cruise track of RV SONNE Cruise SO305 BIOCAT-IIOE2.

3.3.1 Unforeseen Deviations from the Original Agenda of the Cruise

- 1) The cruise was originally planned to take place in spring 2021 but was shifted to April/May 2024 because of the Corona pandemic.
- 2) The originally planned stations in the EEZ of Bangladesh (i.e. on the shelf off Bangladesh) have been cancelled a few days prior to the departure from Colombo because an appropriate diplomatic permission was missing. The berth reserved for an observer from Bangladesh was, therefore, not occupied.
- 3) The national Bangladesh Contact Point for matters relating to the Nagoya Protocol never replied to our repeated requests for approval of collection of biological samples in the EEZ of Bangladesh.
- 4) We left the port of Colombo (Sri Lanka) with a delay of about 3.5 days because the six transport containers did not arrive in Colombo in time. One dangerous goods container is still in Singapore while writing this report (31 July 2024).
- 5) Due to a medical emergency caused by an accident at work, the cruise was terminated prematurely on 11 May. We arrived in Singapore on 16 May, six days earlier as originally planned.

3.3.2 Responsible Marine Research

Sampling and measurements (incl. hydroacoustic surveys) in both the water and in the atmosphere were carried outside the EEZs in international waters only. All drifting instruments (except for the

two Argo floats) were successfully recovered. The deep-sea mooring was deployed in international waters and will be recovered in the upcoming 2-3 years. No zoological material was collected. All data will be archived and will be published after 2 years after the end of the cruise.

4 Narrative of the Cruise

We left the port of Colombo (Sri Lanka) at 20:30 LT on 13 April with a delay of about 3.5 days because the delivery of our six transport containers was delayed. Underway measurements in the surface water and the atmosphere started after leaving the EEZ of Sir Lanka on 15 April at around 05:00 LT (= UTC + 5.5h). After a transit of 2.5 days we performed a test station (stat #01) close to the equator at 0°27.4'N 86°10.7'E on 16 April to test the CTD/Ro and to optimize seawater sampling from the Niskin bottles. The first regular station (stat #02) was performed at 1.5°S 88°40'E on 17 April. It was followed by a series of four stations (stat #03 - #06) on a south to north section crossing the equator along 88°40'E up to 1°N with a latitudinal spacing of 0.5°. At station #04, located at equator (0°N 88°40'E), a long-term deep-sea mooring (with ADCP, oxygen sensors and CTD devices) was deployed in about 4500 m water depth. We then steamed a little further west to 88°E, to the start of our main south to north section through the Bay of Bengal. Stations #07 to #10 were spaced 1° apart from 2°N to 5°N along 88°E. On 21 April we had the first 24h-station (stat #10 at 5°N 88°E). At this station we deployed a first set of drifting particle traps and an Argo float. We continued the section along $88^{\circ}E$ with station #11 (sampled at $6^{\circ}N$ 88°E on 23 April). After completing station #11 we steamed back south to pick up the drifting traps at 5°0.3'N 88°10.8'E (= stat #12 on 23 April I the afternoon). Between 24 April and 26 April, stations #13 (7°N) to #18 (12° N) located in 1° intervals along 88°E were sampled. At station #18, which was the second 24h-station, we deployed both a second set of drifting particle traps and an Argo float. After completing station #18 we followed a diagonal section to the northwest with stations #19 (12°12.3'N 87°1.5'E on 28 April in the morning) and 20# (12°30'N 86°0'E on 28 April in the afternoon). After station #20 we steamed back to pick up the drifting traps at 11°55.7'N 87°43.9'E (= stat #21). We completed the diagonal section with stations #22 (at 12°44.5'N 85°4.5'E on 30 April) and #23 (13°N 84°E on 30 April in the afternoon). This was followed by stations #24 (13°N 85°E on 01 May) to #28 (13°N 89°E on 03 May) which were located in 1° intervals along the west to east section at 13°N. A second diagonal section to the northwest was started with stations #29 (13°30.7'N 88°30.6'E) and #30 (at 14°N 88°E on 04 May). Station #30 was the third 24-station. At this station we deployed the third set of drifting particle traps. The diagonal section was completed with station #31 (14°30'N 87°E on 05 May) and #32 (15°N 85°45'E on 06 May). Station #32 was the fourth 24h-station. After station #32 we steamed back to pick up the drifting traps at 13°59.11'N 88°11.7'E on 07 May (= stat #33). This was followed by stations #34 (15°N 88°E on 08 May) to #37 (15°N 89°36'E on 09 May) which were located along a west to east section at 15°N. Station #37 was the fifth 24h-station. The last station (#38) was performed at 16°N 89°36'E on 11 May. Due to a medical emergency caused by an accident at work on 06 May, the cruise was terminated prematurely at 10h on 11 May after the station work at station #38 had been completed. All underway measurements in the atmosphere and in the water were stopped on 11 May at 14:00 LT (= UTC + 6h) with the entry into the EEZ of India. We arrived at the pier in Singapore at 11:00 LT (= UTC + 8h) on 16 May.

5 Preliminary Results

5.1 Hydrographic Observations

5.1.1 CTD System, Oxygen Measurements and Calibration

(R. Czeschel¹, F. Duerkop¹, H. Großelindemann¹, C. McKellar¹, M. Müller[!],
Q. Zhong¹)
¹ GEOMAR

5.1.1.1 CTD Rosette System

During SO305 BIOCAT-IIOE2 a total of 114 CTD-profiles and 2357 water samples were collected. The rosette system was installed in a Seabird Rosette System frame for 24 bottles. All casts were made with 22 bottles installed. Depth profiles up to a maximum pressure of 4482 dbar were performed. On average, the full water column was sampled at every third stations. Data acquisition was done using Seabird Seasave software version 7.26.7. Pre-processing was done with SBE Data Processing 7.26.7. Some of the Niskin bottles did not close reliably, therefore the corresponding hooks of the water carousel have been replaced and the water carousel has been cleaned. At CTD Station #3 the pump did not work due to a defect conductivity sensor. After we replaced the conductivity sensor the CTD system worked fine at first. During the upcast of CTD Station #19, the pump did not work reliably. After we replaced the connection cable between CTD and conductivity sensor the CTD system worked fine throughout the cruise. Additional to all regular sensors (P, T, S, O) a Chl-fluorescence and turbidity sensor FLNTU manufactured by Wetlabs, a CDOM, and a PAR sensor were attached to the frame of the CTD-rosette system. The cable connection between turbidity sensor and CTD had to be changed after CTD cast #37. Afterwards the turbidity sensor as and all other sensors provided high quality data throughout the cruise. At CTD cast #83 the protecting cap of the turbidity sensor was not removed resulting in unrealistic high data.

The exact configuration of the CTD system can be found in Table 5.1. Additionally, a high precision oxygen sensor (MicroTail), two self-recording LADCPs, a self-recording, self-powered UVP5 (see section 5.6), and a self-recording nutrient sensor (OPUS #71F9) were attached to the water sampler. The spectrometer measure in situ the absorption of UV light by seawater. From comparison with the absorption of clear water and water with a known concentration of nitrate, the nitrate concentration in the seawater sample can be derived. A processing toolbox for OPUS spectrometers had been developed at GEOMAR. The NO_x (NO₂⁻ + NO₃⁻) concentrations resulting from the processing still require a calibration comparable to that of the CTD's conductivity and oxygen sensors.

Processed preliminary CTD data, 5-dbar binned, was sent in near real time to the Coriolis Data Centre in Brest, France, (via email: codata@ifremer.fr) for integration in the databases to be used for operational oceanography applications and the WMO supported GTS/TESAC system.

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	CTD system SBE#9
Pressure sensor	# 0410
T primary	# 5806
T secondary	# 5807
C primary #1	# 3300 (profile 1-3)
C primary #2	# 2512 (profile 4-114)
C secondary	# 4062
O ₂ primary	SBE 43 # 0631
O ₂ secondary	SBE 43 # 2600
PAR Sensor	# 70714
Altimeter	# 42299
Fluorometer/Turbidity sensor	FLNTURTD-2928
CDOM	FLCDRTD-2687

Table 5.1Summary of CTD system SBE #9 configuration and during SO305.

5.1.1.2 CTD Conductivity Calibration

Overall, 158 calibration points were obtained by sampling for salinity. Due to the change of the primary conductivity sensor after profile #3, only 154 calibration points were used for the calibration of the second primary conductivity sensor. Salinity samples were taken by the CTD watch in 'Flensburger' bottles, which have been proven to be ideal for storing salt samples. The measurements with one of GEOMAR's OPTIMARE Precision Salinometers (OPS) are described in section 5.1.2. A simple outlier removal method was applied that discarded the 33% samples with the largest deviations between CTD and bottle samples prior to calibration. The projection of data taken during the bottle stops of the upcast to the data from the downcast was done using the WOCE recommendations by searching within a 30-dbar pressure interval for similar potential temperatures. The conductivity calibration of the downcast data was performed using a 1st order linear fit with respect to temperature, pressure and conductivity (Table 5.2). The calibration results in a salinity RMS-misfit for the downcast of order 0.002145 psu for the second primary and 0.00211 psu for the second ary sensor. The up-cast calibration surpasses these values with a RMS-misfit of 0.001969 psu for the primary and 0.002082 psu for the secondary sensor.

systems used during 50505.	-	-
	CTD system SBE 911	CTD system SBE 911
Sensor pair	Cond. Primary #2512	Cond. Secondary #4062
RMS misfit after calibration - salinity	0.0021447	0.0020858
Polynomial coefficients - conductivity	Offset: 0.046616	Offset: 0.020858
	P1: 9.027e-07	P1: 3.4209e-07
	T1: 0.0016065	T1: 0.00067976
	C1: -0.016658	C1: -0.0073755

 Table 5.2
 End of cruise salinity and pressure summary of downcast calibration information for the two CTD systems used during SO305.

5.1.1.3 Oxygen Calibration

The CTD oxygen downcast for CTD systems is calibrated by using the best 66% of the joint data pairs between downcast CTD sensor value and Winkler-titrated oxygen. For the calibration a correction polynomial depending on pressure, temperature, oxygen and the product of pressure and oxygen was fitted (Table 5.3). A total of 295 oxygen samples were taken. In the end, 196 different oxygen data points were used, which resulted in an RMS-misfit for the downcast of 0.8024 μ mol kg⁻¹ for the primary SBE43 and 0.781234 μ mol kg⁻¹ for the secondary SBE43.

50305.		
	Oxygen Sensor #631	Oxygen Sensor #2600
Sensor pair	primary	secondary
RMS misfit after calibration - oxygen	0.8024	0.78123
Polynomial coefficients - oxygen	Offset: 3.88	Offset: 3.3656
	P1: -0.00037329	P1: -0.00053824
	T1: -0.10493	T1: -0.072756
	O1: -0.078256	O1: -0.08279
	P*O: 1.2599e-5	P*O: 1.1036e-5

 Table 5.3
 End of cruise downcast oxygen summary of calibration information for the CTD system used during SO305.

5.1.2 Conductivity Measurements

(H. Großelindemann¹, C. McKellar¹)

¹ GEOMAR

In order to calibrate the conductivity sensors of the CTD system, the conductivity of 158 water samples was measured using the GEOMAR OPTIMARE Precision Salinometer (OPS) SN 020. Prior to measuring the conductivity with the OPS, the bottle samples have to be degassed to remove gas micro-bubbles as the OPS is sensitive to gas bubbles in probes. Salinity samples were taken with 'Flensburger' bottles from the CTD. Once a box of bottles was full, the bottles were taken into a warm water pool of about 40°C for an hour. Then the bottles were shortly opened to degas. Afterwards the sample bottles were brought to the salinity lab where their conductivity could be measured after 24 hours of cooling down to the lab temperature. The measurement procedure always started by rinsing with old substandard seawater and a test measurement, in order to figure out whether the measurements were stable. This was then followed by the actual standardization with the IAPSO Standard Seawater and a new substandard measurement. Afterwards the measurements of the samples were started. After every tenth sample bottle another substandard was measured. This was done to test whether the instrument is drifting. Mostly the substandard measurements yielded reasonable results. The ending of a measurement series was again followed by another substandard and finalized by re-measuring the remaining Standard Seawater from the standardization in the beginning, which was expected to be slightly higher (0.001-0.002) than the original standard measurement.

In addition to the salinity samples from the CTD, 44 samples from the thermosalinograph (TSG, see section 5.1.3) were also measured. These were used for the calibration of the TSG. Furthermore, a couple of samples were measured for other groups (GoFlo and sediment traps), because they needed precise salinity values for their own investigations.

5.1.3 Thermosalinograph

(H. Goßelindemann¹, C. McKellar¹)

¹ GEOMAR

Underway measurements of sea surface temperature (SST) and sea surface salinity (SSS) were continuously done by the ship's dual thermosalinograph. One inlet is located at the portside (TSG1) while the other thermosalinograph's inlet is at the starboard side (TSG2). The parallel system worked well throughout SO305. Due to the switch off of scientific measurements in the EEZ of Sri Lanka and before entering the EEZ of India for the transit to Singapore there are no measurements for these areas either. A total of 44 salinity samples were taken during the cruise and measured during the transit with the GEOMAR OPTIMARE Precision Salinometer (OPS) SN 020 for calibration.

5.1.4 Underway CTD (uCTD)

(R. Czeschel¹, F. Duerkop¹, H. Großelindemann¹, C. McKellar¹, M. Müller¹, Q. Zhong¹)

¹ GEOMAR

The Underway CTD (uCTD) of Oceanscience was used for underway profiling of upper ocean temperature, salinity, and pressure. The system consisted of a Sea-Bird CTD instrument with embedded data acquisition. The sensor is contained within a streamlined pressure case with an integrated line spool. The winch holds 1400 m of Spectra line for profiles to over 400 m depth. The tail rewinder loads the tail spool with the Spectra line. The uCTD was installed on the back of the working deck. The nominal operating depth of the uCTD depends on the vessel speed. We used the uCTD while steaming with ~8kn. The profile depth was up to 250 m depth. We used the uCTD to measure between our CTD stations between 5°N and 7°N while we were crossing an eddy. In total, 15 profiles were recorded with the uCTD (see Table 11.1 in the Appendix). After two profiles the laser of the Rewinder displayed an error. After cleaning the light barrier, the uCTD worked again. After another profile, a pin which was used to lock the tail in the rewinder when winding up the Spectra line on the tail got stuck. The pin was then replaced by a screw. From then on, the uCTD worked well without any problems and another 12 profiles were recorded. We stopped recording after we had crossed the eddy.

5.2 Current Observations

5.2.1 Vessel-Mounted ADCP

(R. Czeschel¹)

¹ GEOMAR

Underway-current measurements of the upper ocean were performed continuously throughout the entire cruise (except in the territorial waters of Sri Lanka, India, Myanmar and Singapore) using two vessel-mounted Acoustic Doppler Current Profilers (vmADCP): a 75kHz RDI Ocean

Surveyor (OS75) and a 38kHz RDI Ocean Surveyor (OS38) both mounted in the ship's hull. Measurements started at 2.9°N, 83.11°E on 15 April 2024, 02:38 UTC. We stopped the record of both vmADCP at the EEZ of India at 15.17°N, 89.63°E on 11 May 2024, 08:00 UTC. Due to the test of the hydrophone for the mooring deployment we stopped recording of vmADCP data on 17 April from 07:49 to 08:59 UTC. The 75 kHz ADCP broke down twice after stop/start recording. Therefore, the following files are corrupt:

- 1) so305os75004_000000.ENX, *ENR, *LOG, *N2R, *N3R, *NMS, *VMO so305os75005_000000. ENX, *ENR, *LOG, *N2R, *N3R, *NMS, *VMO
- 2) so305os75013_000000.ENX, *ENR, *LOG, *N2R, *N3R, *NMS, *VMO

After the vmADCP was restarted, the data was accidently saved under the file name of the previous cruise. Therefore, following files should be renamed:

- 1) so304_os75_304_00000* -> so305os75005_00000*
- 2) so304_os75_305_00000* -> so305os75006_00000*

The OS75 and the OS38 are aligned to zero degrees (relative to the ship's center line). Both instruments ran in narrowband mode. The OS75 instrument was configured with 100 bins of 8 m and a blanking distance of 4 m, pinging 24 times per minute and reaching a range of 600 m to 700 m. The OS38 used 55 bins of 32 m and a blanking distance of 16 m, pinging 19 times per minute and reaching a range between 1200 m and 1500 m. During the entire cruise, the SEAPATH navigation data was of high quality. No interference with the 12kHz echosounder EM122 that delivered high quality bathymetry data was detected. Post processing of the data was carried out separately for each instrument. The applied mean misalignment angles and amplitude factors with the associated standard deviation are summarized in Table 5.4.

DIE 5.4		vessei-mo	Sumed ADCP cambration.	
	OS	Mode	Misalignment angle \pm std	Amplitude factor \pm std
	75	NB	$-0.1181^{\circ} \pm 0.5368^{\circ}$	1.0007 ± 0.0094
	38	NB	$-0.2790^{\circ} \pm 0.6262^{\circ}$	1.0049 ± 0.0136

Table 5.4Vessel-mounted ADCP calibration.

5.2.2 Lowered ADCP

(R. Czeschel¹, F. Duerkop¹, H. Großelindemann¹, C. McKellar¹, M. Müller¹, Q. Zhong¹)

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<sup>1</sup> GEOMAR
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During the whole cruise the CTD/Rosette system was equipped with a lowered ADCP setup based on two Teledyne RDI ADCPs. The setup consisted of an upward looking and a downward looking 300-kHz instrument. These two instruments were mounted inside the CTD rosette with specially manufactured frames protecting the instruments and allowing zero obstruction of the acoustic beams. The LADCP system worked without trouble with SN #11461 as downward-looking master instrument and #11436 as upward-looking slave during the whole cruise. During the cruise we used a software, which controlled the start, stop, download, and erase of the cycles of the two LADCP systems (ladcp_tool_1.9.3 developed at GEOMAR). An energy supply system that draws energy for the ADCPs from the CTD system using rechargeable capacitor (10.000µF / 100V)

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worked well throughout the cruise. The start as well as the download signal for the LADCP system was conducted via Bluetooth. The distance between the LADCP instruments and the Bluetooth transmitter was reduced by installing the transmitter at a tension belt that was stretched in the hangar next to the CTD to minimize communication problems. Therefore, the communication with the LADCPs via Bluetooth worked well throughout the cruise. Data processing took place during the cruise using the GEOMAR LADCP processing software V10.22, which includes both shear and inversion methods to derive an absolute velocity profile. As additional data are necessary for the processing, the corresponding pre-processed CTD files were used containing pressure, temperature and salinity profiles as well as time and navigation data. The instruments of Teledyne RDI instruments delivered very good deep-ocean velocity profiles when processed in conjunction with the observations of the vmADCP and when coming close enough to the seafloor to obtain TRDI bottom track data.

5.3 Floats

(M. Müller¹, R. Czeschel¹)

¹ GEOMAR

During SO305, two NKE floats (= Argo floats) were deployed (Table 5.5). These profiling floats are measuring temperature, salinity, oxygen and pressure of the upper 2000 m of the ocean. The Argo floats were tested, activated and then deployed by hand.

Table 5.5Float deployments during SO305.

Tuble 2.2 Thou deployments during 50505.					
Float serial number	WMO	Latitude	Longitude	Deployment Date (UTC)	
AI2600-22DE009	6904212	11°20.222'S	012°14.944'E	26-Apr-2022 01:26	
AI2600-22DE008	6904213	11°10.467'S	011° 13.183'E	26-Apr-2022 14:41	

5.4 Mooring

5.4.1 Mooring Operations

(M. Müller¹, R. Czeschel¹, F. Duerkop¹)

¹ GEOMAR

Mooring work during SO305 BIOCAT-IIOE2 consisted of the deployment of a long-term mooring at the equator at 89°E (Table 5.6). This mooring consists of a WH-ADCP (up-looking) and a Signature-ADCP (down-looking) measuring the strength of the upper currents as well as oxygen (Optodes), temperature and salinity (MicroCats), McLane moored profiler (849 - 3324m) and single point current meters (Aquadops) just below the ADCP to analyse water mass variability.

Mooring Deploy	ment				Notes:	KPO_1234
Vessel:	Sonne	SO305				
Deployed:	18-April	2024	12:46 UTC			
Vessel:						
Recovered:						
Latitude:		00°	01.461'	S		
Longitude:		88°	40.076'	Е		
Water depth:		4513	Mag Var:	2.2		
	Designed					
ID	Depth	Instr. Type	s/n	Start-up	Remarks	
KPO_1234_01	100	XEOS-XMA Argos	214435	ready		
KPO_1234_02	100	Float	Ani 4			
KPO_1234_03	100	WH ADCP 300kHz	2379	X		
KPO_1234_04	100	Signature ADCP	200065	X		
KPO_1234_05	199	Microcat	3196	X		
KPO_1234_06	199	O2 Logger (ind. Opt.)	054	X		
KPO_1234_07	303	Microcat	8947	X		
KPO_1234_08	303	O2 Logger (ind. Opt.)	101	Х		
KPO_1234_09	503	Microcat /p	6855	Х		
KPO_1234_10	503	O2 Logger (ind. Opt.)	078	X		
KPO_1234_11	849					
	3324	MMP	14754	Х		
KPO_1234_13	3631	Aquadopp down /p	16454	X		
KPO_1234_14	3942	Aquadopp down /p	16453	X		
KPO_1234_15	4252	Aquadopp down /p	16456	X		
KPO_1234_16	4368	Microcat /p	6859	х		
				Mode:		
KPO_1234_17	4504	Release AR661	642	А	Enable: 4A8	33 / Release: 4A84
KPO_1234_18	4504	Release AR861	1645	Mode: B	Enable: 0A8	BA / Release: 0A55

Table 5.6Mooring operations during SO305.

5.4.2 Calibration of Moored Instruments

(M. Müller¹, F. Duerkop¹, R. Czeschel¹)

¹ GEOMAR

CTD/O₂ cast calibrations were performed for all MicroCATs, Optode loggers and Aquadops as pre-deployment calibration (CTD casts 2 and 4) by attaching the instruments to the CTD frame. During the upcast, 5 and 3 calibration stops were done over the whole profile range (depths chosen at low gradient-regimes for the respective parameters). Each stop had a duration of at least 6 min in order to ensure fully adjusted measurements at the calibration points. Additionally, releaser tests were performed at CTD cast 2. Lab calibrations at GEOMAR were conducted for all O₂ loggers in water-filled beakers of 0% and 100% O₂-saturated water at two different temperatures (~6°C and ~21°C) following the Aanderaa Optode manual.

5.5 Shipboard Microstructure Measurements

(R. Czeschel¹, F. Duerkop¹, H. Großelindemann¹, C. McKellar¹, M. Müller¹, Q. Zhong¹)
¹ GEOMAR

A microstructure (MSS90) profiler (#028) of Sea and Sun Technology was used to infer turbulent dissipation rate and diapycnal diffusivity, aimed at calculating turbulent fluxes of oxygen, heat, momentum, nutrients, and nitrous oxide (N₂O). The loosely tethered profilers are equipped with 2 airfoil shear sensors (#121, #122) and a fast thermistor, as well as some common CTD sensors: pressure, conductivity, temperature and turbidity sensor. In total, 120 profiles were recorded on 42 MSS stations to a maximum depth of 320 m. Most stations consisted of 3 microstructure profiles between 2 CTD casts. A list of all MSS profiles is given in Table 11.2 in the Appendix.

The MSS worked fine in the beginning of the cruise. However, at profile #55 the MSS started to have some data dropouts. In the following stations the dropouts got more. The analyse with the time domain reflectometer (TDR) didn't show any clear hints of a problem with the underwater cable. Since the most likely problem should have been in the first few meters of the cable, the cable was shortened about 10 m before profile #81. The following profiles still had some severe data dropouts that's why the whole underwater cable was exchanged before profile #85. Again, there was not much improvement in the following profiles. Before profile #101 the MSS itself was opened to look for any water within the device. No signs of water were found but two dry bags were added to prevent condensation water. The IC of the RS485 interface was exchanged. Also, all connectors have been thoroughly cleaned and visually checked. An insulation measurement of the cable didn't show any problems. In the following profiles the data dropouts were less, but still not completely gone. Often the first profile had the most dropouts and the data dropouts of the following profiles decreased for some reason. The dropouts appeared to come up randomly, sometimes on the up- and sometimes on the down cast. They also came up at different depth. The brushes of the slip rings might be worn out, but that couldn't be verified on board due to the high risk of losing some parts while doing that.

The fall rate of the profilers was 0.55 m/s in the beginning. Before profile #101 additional weight (2 thin rings) was added in the hope to bring the high noise level (~1 10^{-8} m²/s³) down. That increased the fall rate to about 0.85 m/s. As the noise level was still high, but now the fall rate too high the weight was reduced by each one 1 thin ring before profile #104 (#112, #118) leading to a fall rate of 0.8 m/s (0.7 m/s, 0.55 m/s).

5.6 Underwater Vision Profiler

(M. Müller¹, R. Kiko¹)

¹ GEOMAR

During all regular CTD casts, an Underwater Vision Profiler 5 HD (UVP5 HD; serial number 28) was operated on the CTD rosette. The instrument consists of one down-facing HD camera in a 6000 dbar pressure-proof case and two red LED lights which illuminate a 1.24 L water volume. During the downcast, the UVP5 takes 20 pictures of the illuminated field per second. For each picture, the number and size of particles are counted and stored for later data analysis. Furthermore,

images of particles with a size > 500 μ m are saved as a separate "Vignettes" - small cut-outs of the original picture – which allow for later, computer-assisted identification of these particles and their grouping into different particle, phyto- and zooplankton classes. Since the UVP5 was integrated in the CTD rosette and interfaced with the CTD sensors, fine-scale vertical distribution of particles and major planktonic groups can be related to environmental data. In total 90 UVP5 profiles could be obtained. At each station with a water depth < 6000 m a full-depth profile was obtained. Further, computer-assisted analysis of the approximately 700,000 images taken with the UVP5 will be done in the home laboratory in order to reveal fine-scale distribution patterns of particles and zooplankton.

5.7 Dissolved Nutrients and Dissolved Oxygen

(L. Schmidt¹, J. Usinger², K. Lange², H. Bange²)

¹ Hereon, ² GEOMAR

5.7.1 Methods

Dissolved inorganic nutrients (nitrate, nitrite, silicate and phosphate) were measured with a continuous flow analyser (CFA) according to the photometric methods described in Grasshoff et al. (1999): The right amount of reagent is added to the sample and mixed to form a colour complex. Its concentration is quantified by the build-in photometer. In contrast to the other nutrients, nitrate is determined by reducing the nitrate to nitrite with cadmium. The nitrate concentration is calculated by subtracting the nitrate-derived nitrite concentration from the total concentrations of nitrite. The CFA works automatically and needs less than 2 mL of sample for the determination of all four nutrients. Approximately 50 samples per hour can be measured.

Dissolved ammonium was analysed with the ortho-phthalaldehyde (OPA) fluorometric method following Holmes et al. (1999). The working reagent containing OPA is forming a fluorescent complex with ammonium. It can be measured within 12 to 24 hours. The efficiency of the OPA is dependent of the matrix of the sample. Therefore, low nutrient sea water from a recent station is used for the calibration using the method of standard addition.

The Winkler method was used to determine dissolved oxygen. The sample was taken air free for all stations. Then manganese chloride and alkaline sodium iodide were simultaneously added to the sample. Manganese is oxidized by the dissolved oxygen and forms a precipitate. By adding sulfuric acid, the oxidized manganese dissolved and converted iodide to iodine which resulted in a yellow colour. Then, iodine was titrated with sodium thiosulfate until the solution almost lost its colour. Zinc iodide starch solution was added and coloured the solution in purple. The titration continued until the solution reached transparency. The titrated volume was used to calculate the oxygen concentration via: O_2 (µmol/L) = a * f * 5000 / (b-2), with a representing the titrated volume (in mL), b the volume of the sampling flask and f the factor of the sodium thiosulfate solution.

Reusing the lids of the sampling vials would result in too high nitrate concentrations if they were not rinsed properly while sampling. As the ammonium concentrations were very low, avoiding contamination and achieving good replicates was difficult.

Bubbles were only present in oxygen samples in near-surface waters. Near-surface waters are more than 10 °C warmer than the room temperature. While cooling down, the shrinking volume might have caused the bottle to suck in air from outside. These bubbles might have influenced the measurements. The lids of the glass bottles are prone to slipping out easily, so it is advised to use clippers to prevent accidentally opening the bottle.

All measurements and quality control were finished during the cruise. No samples will be transported to GEOMAR, and all data is already processed.

5.7.2 Sampling

All dissolved nutrients and dissolved oxygen were sampled from every CTD station. Furthermore, the CFA measured nutrients were sampled and measured for the underway system, the GoFlo bottle sampling, and the Towed-fish. Besides samples of the two on-board running incubations were taken and nutrients were analysed. Sampling from the underway system was organized by Dennis Booge, GEOMAR. Nutrient triplicates were taken every three hours. The underway system is sampling surface water from 5 m water depth. The GoFlo and towed-fish sampling was under the responsibility of Martha Gledhill, GEOMAR. Dennis Booge and the participants from the SDU were running and sampling their incubations.

At the first CTD station, the nutrients were sampled as triplicates to check the standard deviation and necessity of taking triplicates. The following CTD casts were sampled as single sample. Ammonium was always sampled separately as triplicates. At the first stations, one replicate was taken for every depth. After further evaluation, triplicates were taken at two specific depths to determine the accuracy of the measurement. The chosen depths were at the oxygen minimum and maximum. From station 5 onwards, the sampling procedure was changed to sampling triplicates at previously specified depths including bottom, oxygen maximum, oxygen minimum, chlorophyll maximum, and salinity minimum. This reduced the number of samples which needed to be measured and increased the accuracy of the calibration for the O_2 sensor of the CTD. It was avoided to sample too close to the surface. As the oxygen concentration strongly changes close to the chlorophyll maximum and, thus, provides unreliable data for calibrating the O_2 sensor of the CTD, sampling this depth was discarded later on.

5.7.3 Preliminary Results

At 0°59.969' S and 88°40.209'E (Station 2), the oxygen minimum zone was not yet reached. Below the fully oxygenated surface, the oxygen concentrations declined to approximately 50 μ mol/L in the depth range below the euphotic zone till 1000 m including a secondary oxygen maximum in 430 m with 78.44 μ mol/L. Below 1000 m, oxygen concentrations were increasing with depth. Nitrite was only present between 50 and 100 m with concentrations lower than 0.96 μ mol/L. Ammonium concentrations fluctuated between 0.04 and 0.20 μ mol/L. All other nutrients were depleted in the surface and increased with depth. At depth, the nutrients showed maximal concentrations of approximately, 36.83 μ mol/L for nitrate, 2.43 μ mol/L for phosphate and 134.36 μ mol/L for silicate (Figure 5.1).

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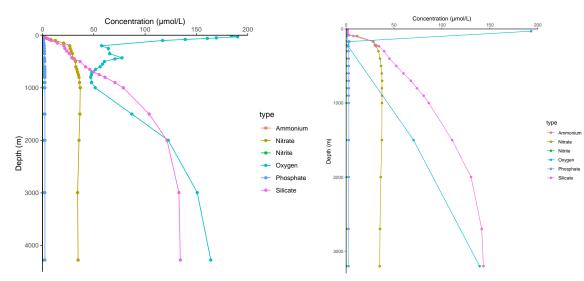
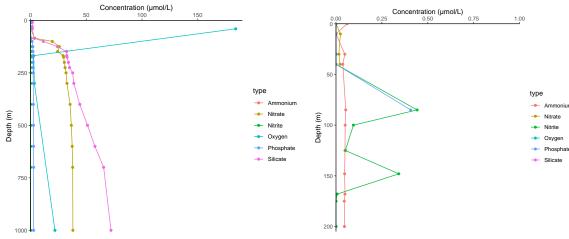
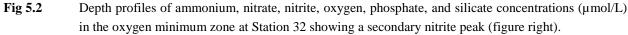


Fig 5.1 Depth profiles of ammonium, nitrate, nitrite, oxygen, phosphate, and silicate concentrations (in µmol/L) outside the oxygen minimum zone (figure left: station 2) and in the oxygen minimum zone (figure, right: station 24)

At 13°00.097' N and 85°00.062'E (station 24), low oxygen concentrations were found. The surface was fully oxygenated. Minimal oxygen concentrations of 2.08 μ mol/L were reached at 233 m depth. Below the oxygen minimum zone, oxygen concentrations were increasing with depth. Nitrite was only present between 50 and 100 m and at maximum depth with a maximum concentration of 0.41 μ mol/L in 83 m. Ammonium concentrations fluctuated between 0.03 and 0.05 μ mol/L. All other nutrients were depleted in the surface and increased with depth. At depth, the nutrients showed maximal concentrations of approximately, 37.59 μ mol/L for nitrate, 2.86 μ mol/L for phosphate and 143.23 μ mol/L for silicate (Figure 5.1).





At 14°58.149' N and 85°43.914'E (station 32, cast 88 and 91), the lowest oxygen concentration was 2.80 μ mol/L in 168 m. Again, the surface was fully oxygenated, and oxygen concentrations were increasing with depth below the oxygen minimum zone. Nitrite showed a regular maximum at 85 m with 0.44 μ mol/L and a secondary nitrite peak at 148 m with 0.34 μ mol/L. Ammonium concentrations fluctuated between 0.04 and 0.06 μ mol/L. All other nutrients were depleted in the

surface and increased with depth. At depth, the nutrients showed maximal concentrations of approximately, $38.04 \mu mol/L$ for nitrate, $2.82 \mu mol/L$ for phosphate and $72.19 \mu mol/L$ for silicate within the first 1000 m (Figure 5.2, data referring to station 32, cast 91).

5.8 Nitrous oxide and Methane

(I. Hentschel¹, L. van Bonn¹, I. Stoltenberg¹, H. Bange¹)

¹ GEOMAR

5.8.1 Introduction

The major goal of the working group nitrous oxide and methane during the SO305 BIOCAT-IIOE2 cruise was the determination of dissolved nitrous oxide (N_2O) and methane (CH₄) throughout the water column in the Bay of Bengal. Both gases are microbially produced in the water column and represent the second and first most important greenhouse gases, respectively. The global warming potential (GWP100) of methane equals 27.9. Thus, one methane molecule has the same radiation force as 28 molecules of CO_2 (Pachauri et al., 2014). The GWP100 for nitrous oxide is about 298. Most of the atmospheric methane is formed via biological production called methanogenesis. It is typically believed that the major formation of methane is performed under strictly anoxic conditions, however, several processes have been suggested to produce CH₄ under oxic conditions as well (Bange et al., 2024). Nitrous oxide is formed dominantly as a byproduct via nitrification under oxic/suboxic conditions in the water column. Its formation increases under suboxic/anoxic conditions as an intermediate product of denitrification. Under anoxic conditions N2O is consumed via denitrification. Therefore, the enrichment of nitrous oxide in the water column depends on the oxygen concentration (Bakker et al., 2014). In view of the ongoing climate change, it is essential to understand the global natural and anthropogenic pathways of the two greenhouse gases.

5.8.2 Methods

Samples for methane and nitrous oxide were collected with special focus on the oxygen minimum zone and the surface water. Nitrous oxide samples were measured on board using static headspace analysis. For this, brown 20 mL vials were filled air bubble-free directly from the Niskin bottles at the CTD/RO and sealed with rubber stoppers (Butylgummihohlstopfen, Chromatographiehandel Müller, Fridolfing, Germany) and aluminium caps (Bördelkappe R20-oA, Chromatographiehandel Müller, Fridolfing, Germany). All the samples were immediately poisoned with 50 μ L of saturated aqueous mercury chloride (HgCl₂) solution. Afterwards, a headspace of 10 mL Helium (He 5.0, Air Liquide GmbH, Düsseldorf) was injected into the sample vial with a gas-tight syringe (Series A-2 Syringe 10 mL, VICI Precision Sampling, Baton Rouge, LA, USA). An empty 20 mL syringe (20mL BD PlastikpakTM, BD Heidelberg, Germany) was injected through the rubber stopper of the sample vial beforehand for pressure compensation. This method was applied to every sample taken. The samples were vortexed for 30 seconds using a vortex shaker (Vortex Genie 2, Scientific Industries Inc., New York, USA) and then stored for at least two hours. A solubility equilibrium develops between the liquid phase and the gas phase (= headspace), as the dissolved gas from the sample diffuses into the overlying gas phase. After two

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hours the equilibrium state is reached, and the samples can be measured. For this, 9 mL of the headspace was transferred to a 10 mL gas-tight syringe. Any anomalies that may have occurred up to this point, such as inconsistencies in the amount of compensation fluid or residual liquid when extracting the headspace, were recorded in the measurement protocol. The sample was injected into the gas chromatograph (Hewlett Packard HP 5890 Series II, Agilent Technologies, Santa Clara, California, USA) via a two-position valve (Valco Valve, VICI International, Schenkon, Switzerland). It passes through a dry trap filled with phosphorus pentoxide (P₂O₅, Sicapent[®], Merck, Darmstadt, Germany) before entering the sample loop. By switching the valve, 2 mL aliquot were injected onto the column together with the carrier gas (argon/methane gas mixture 95/5, AirLiquide GmbH, Düsseldorf, Germany). The column used is a 6' 1/8" stainless steel column packed with a 5 Å molecular sieve (Grace GmbH & Co. KG, Worms, Germany), which effectively separates nitrous oxide. The GC is equipped with an electron capture detector (ECD) which was operated at 350°C. The GC temperature was set to 190°C. The argon/methane was set to a flow rate of 30 mL min⁻¹. Samples were run every 2.40 minutes and every 9.40 minutes. Before sampling, the GC was calibrated daily. Therefore, two standard gas mixtures with different dilutions were injected in triplicates and a calibration curve with a quadratic fit was applied. A correlation coefficient of R²>0.95 had to be met. The two standard gases had calibrated concentrations of 330 ppb to 990 ppb of N₂O (Deuste Steiniger GmbH, Mülheim, Germany). By diluting the standard with Helium at atmospheric pressure, a 66% dilution of the 990 ppb standard gas and a 44% dilution of the 330 ppb standard gas mixture was used. In some cases, the sample concentrations exceeded the range of the calibration, then the linear fit was applied. The signals measured at the ECD were further processed with the ChromStar software (ChromStar Version 6.3, Software für Chromatographie und Prozessanalytik GmbH, Wehye-Leeste, Germany). N₂O peaks were integrated manually and transferred to a Microsoft Excel spreadsheet, calculating the final concentrations and standard deviation. During the measurements two dry traps broke due to mechanical pressure, which was then prevented by adjusting the setup at the GC and an additional dry trap was lost due to an excess amount of sample water that had entered the syringe. The calibrations of specific stations vary from the remaining calibrations due to presumably leaking septa at the helium outlet and GC injection port. Upon identifying the source of the error, these were replaced and thereafter renewed every other day.

The methane samples were only poisoned and then stored for measurement in the lab. This will be done by using a purge and trap unit. The purge and trap method relies on continuously passing a carrier gas through the sample, causing the volatile gases in the sample to be released with the carrier gas and collected in a trap stored in liquid nitrogen.

Since the measurement of nitrous oxide data was already completed during the cruise and the data could be collected, the evaluation of these will likely take place a few months after the cruise ends. Predicting the evaluation of methane is difficult, as ongoing issues with the purge and trap unit can lead to time delays, and the method is generally time-consuming. Results can be expected at the earliest during the next year.

5.8.3 Overview Sampling and Measurements

During the SO305 BIOCAT-IIOE2 cruise, we took water samples for methane and nitrous gas from every second station except stations 37 and 38, where we sampled both due to the premature

termination of the cruise. At the 24h-stations, only one depth profile was sampled (stations 10, 18, 30, 32, 37). Additionally, a gradient pump system was applied during sampling from the Zodiac to achieve high-resolution measurements in 0.25 m steps from the sea surface down to a water depth of 2 m. We collected data from the Zodiac for all 24h-stations except station 32 due to bad weather conditions. From regular CTD/Ro profiles, we collected trace gas water samples from the ocean bottom to 1000 m in 1000 m steps. At water depth <1000m, the sampling was continued in 100 m steps for nitrous oxide and 200 m steps for methane. An extra set of samples (triplicates) for both gases was taken if there was a chlorophyll maximum (or two), and oxygen minimum or maximum. Thus, at water depths <100 m we usually had up to 4 samples (e.g. 60 m, 30 m, 10 m, and surface) depending on the CTD profile. All nitrous oxide water samples were measured on board (around 1000 samples). Methane samples will be measured at the GEOMAR in Kiel after the cruise.

5.8.4 **Preliminary results**

The primary results of the measurement show a correlation between oxygen concentrations in the water column and nitrous oxide concentrations. As we descended deeper into the oxygen minimum zone, and oxygen levels more and more decreased, the N₂O concentrations simultaneously increased. They reached their maximum at station 30 with 133.2 μ mol/L. Particularly notable was the observation of the decline in N₂O levels at the depth of the oxygen minimum at station 32, which may be attributable to denitrification processes.

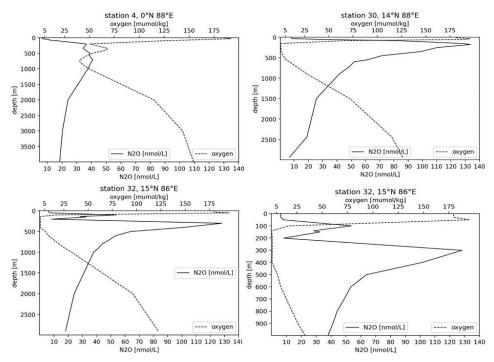


Fig 5.3 Depth profiles of N₂O (in nmol/L and oxygen (in μmol/L) concentrations on the equator (upper left) and at 14°N (upper right) on the 88°E transect and a possible denitrification peak at 15°N 86°E on a full depth profile (lower left) and for the first 1000 m (lower right).

5.9 **Underway Trace Gas Measurements**

(P. Eisnecker¹, M. Sommer¹, D. Arévalo-Martínez^{1,2}, H. Bange¹)

¹ GEOMAR, ² Radboud Univ., Nijmegen, NL

Continuous measurements of dissolved N2O, CO2, CO, and CH4 in seawater were carried out during SO305 BIOCAT-IIOE2 by means of an autonomous equilibrator headspace setup coupled to an off-axis integrated cavity output spectroscopy analyser (LGR; N₂O and CO) and a cavity ringdown spectroscopy analyser (PICARRO; CO2 and CH4) (Arévalo-Martínez et al., 2013). Water was drawn into the system at ~ 6 L min⁻¹ by a LOWARA submersible pump installed in the ship's moonpool (~ 6 m depth). Ambient air measurements were carried out every six hours by drawing air into the system from an air inlet located at deck 10 about 32 m above sea level. Control measurements were performed every 6 hours by means of three standard gas mixtures (DEUSTE STEININGER GmbH) bracketing the expected concentrations in this area. These gas mixtures were calibrated against primary NOAA standards at the Chemical Oceanography Department of GEOMAR Helmholtz Centre for Ocean Research, Kiel.

5.10 **Carbon Monoxide**

(P. Eisnecker¹, M. Sommer¹, H. Bange¹)

¹ GEOMAR

The air-sea exchange and oceanic cycling of climate forcers (or greenhouse gases, GHG), including the indirect GHG carbon monoxide (CO) are fundamental in controlling the evolution of the Earth's atmospheric chemistry and climate. Hence, the investigation of their distribution and sea-air fluxes is pivotal for better understanding potential responses of the ocean and the overlying atmosphere to environmental changes such as warming and deoxygenation. It is wellestablished now that the ocean under the current conditions is a minor source for CO. However, oceanic emission estimates are still associated with a high degree of uncertainty and there are still large knowledge gaps in the processes controlling the production, distribution, and reduction of CO in the ocean (Bange et al., 2024).

Water samples from CTD/Ro stations have partly been taken from different CTD casts, with the primary cast taking deep water samples and the following cast taking mid-depth and shallow water samples. Aimed sampling depths are 100 m, 70 m, 50 m, 30 m, 10 m, surface. Focus depths are adapted or added for peaks in oxygen concentration and the primary chlorophyll α maximum. Triplicates have been taken from each depth. CO concentrations in the samples have been measured immediately on board with a CO analyser (ta3000R SP, Gas Monitor CO in Headspace, AMETEK). Discrete CTD samples from selected CTD stations including the 24h-stations (marked with x) are listed in Table 5.7. Carbon monoxide samples from the surface gradient pump system were taken at 200 cm, 175 cm, 150 cm, 125 cm, 100 cm, 75 cm, 50 cm, and 25 cm water depth while sampling with the Zodiac.

DShip Station ID	24h-Station	Sampled depths [m]
SO305_2_5	-	200, 100, 80, 60, 50, 30, 10
SO305_4_2	-	200, 150, 100, 80, 60, 50
SO305_4_3	-	30, 10
SO305_6_3	-	200, 170, 100, 70, 50, 30, 10
SO305_8_3	-	215, 100, 80, 55, 30, 5
SO305_10_5	Х	90, 70, 43, 30, 10, 2
SO305_10_8	Х	100, 70, 60, 50, 30, 10, 2
SO305_10_12	Х	100, 85, 70, 50, 30, 10, 2
SO305_10_16	Х	100, 85, 70, 55, 30, 2
SO305_13_4	-	150, 100, 75, 60, 50, 30, 10, 2
SO305_15_3	-	100, 80, 65, 50, 30, 10, 2
SO305_18_4	Х	100, 80, 73, 50, 30, 10, 2
SO305_18_9	Х	100, 80, 70, 50, 30, 10, 2
SO305_18_13	Х	100, 80, 70, 50, 30, 10, 2
SO305_18_16	Х	100, 80, 70, 50, 30, 10, 2
SO305_20_3	-	100, 80, 50, 30, 10, 2
SO305_23_3	-	90, 80, 55, 30, 10, 2
SO305_25_3	-	100, 80, 70, 50, 30, 10, 2
SO305_28_2	-	100, 80, 50, 30, 10, 2
SO305_30_3	Х	100, 75, 50, 30, 10, 2
SO305_30_5	Х	100, 75, 50, 30, 10, 2
SO305_30_8	Х	100, 80, 50, 30, 10, 2
SO305_30_10	Х	100, 75, 50, 30, 10, 2
SO305_32_5	X	100, 90, 50, 30, 10, 2
SO305_32_6	X	100, 85, 45, 30, 10, 2
SO305_32_14	X	100, 85, 40, 30, 10, 5
SO305_35_4	-	100, 85, 55, 30, 10, 5
SO305_37_3	Х	100, 70, 40, 30, 10, 3
SO305_37_8	X	100, 60, 50, 30, 10, 2
SO305_37_12	Х	100, 80, 50, 45, 10, 2

 Tab. 5.7
 Station number and sampled depths for carbon monoxide measurements.

5.11 Nitric Oxide (NO) Measurements

(R. Ingeniero¹, H. Bange¹)

¹ GEOMAR

5.11.1 Introduction

Nitric oxide (NO) is a short-lived climate forcer that indirectly contributes to greenhouse gas effects by interacting with other compounds to form ozone, methane, and nitrate aerosols, leading to negative radiative forcing (IPCC, 2021). The determination of dissolved NO concentration is challenging because of its reactivity, which results in a very short lifetime in (sea)water, ranging from 3 to 100 s. Until now, little is known about the distribution as well as the production and consumption processes of NO in the marine environment. Despite these challenges, there has been a growing interest in dissolved NO measurements in different aquatic systems, including open seas, coastal waters, and rivers, over the past decade. Most of the studies have indicated that the marine environment is a source of atmospheric NO. NO is a short-lived intermediate in the

microbial nitrogen cycle (Kuypers et al., 2018), particularly in nitrification, denitrification, and anammox. To this end, this study has two main objectives:

- To estimate the sources and sinks of NO in the ocean/atmosphere. To achieve this, we measured the NO concentration in the near-surface water along the cruise track. Furthermore, using the atmospheric NO concentration measured by the Leibniz Institute for Tropospheric Research (TROPOS), we will estimate the seawater-atmosphere flux densities.
- 2) To determine NO concentration/distribution along the water column in selected stations. Given that the Bay of Bengal is also known as an oceanic region with a pronounced minimum zone, we aim to understand how deoxygenation influences nitrogen cycling processes and, ultimately, how it could influence NO distribution.

5.11.2 Methods

5.11.2.1 Sampling and measurement of NO from the underway and submersible pump

Triplicate bubble-free water samples from approximately 6 m depth were collected from the vessel's underway pump every 6 hours along the cruise track. At the 24h-stations, a submersible sampling pump with a depth sensor was deployed up to approximately 140 meters depth to collect water samples. Approximately >300 samples were measured on board. To ensure that the water samples are taken at and representative of the desired sampling depth, sampling was done 10 minutes after reaching the determined depth. A portable NO calibration source (2BTech Model 714 NO₂/NO/O₃ Calibration SourceTM) was used to calibrate the NO detector (Birks et al., 2020). The resulting gas output from the calibrator covered the detection range of the NO detector from 0 to 1000 ppb NO. NO signal outputs by the NO detector were recorded using PuTTY 0.78, a free and open-source client application for Windows. To determine NO mole fractions, the Rieman integrals of the signal peaks were calculated using the MATLAB (2022b) trapezoidal numerical integration function trapz. Since stripping measurement time was logged manually, it will take some time to process and analyse all the data obtained from the SO305 cruise. This will be done back at GEOMAR and may take approximately 2 to 3 months.

5.11.2.2 Incubation Experiments

A conceptual experiment was conducted to investigate the influence of nitrite concentration on dissolved nitric oxide concentration in seawater. Surface seawater samples were collected using a CTD/Ro sampler and transferred into 50 mL glass vials under bubble-free conditions. The samples were divided into control and treatment groups, with the latter spiked with varying concentrations of nitrite. These samples were incubated under alternating light/dark cycles and complete darkness for 24 and 48 hours. Dissolved NO concentrations were subsequently measured, providing insights into the interaction between nitrite levels and NO production in marine environments.

5.11.3 Preliminary Results

Shown below are profiles of dissolved NO and oxygen concentrations (Fig. 5.4). The NO concentration seems to increase in the area with low oxygen concentration, probably due to microbial NO production from the denitrification process.

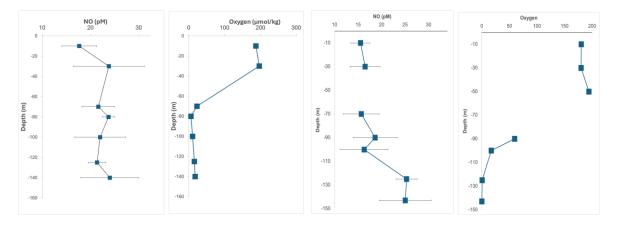


Fig 5.4 Depth profiles of NO (pM) and oxygen (μmol kg⁻¹) concentrations at the 24h-station on 27 April 2024 (figure left) and 6 May 2024 (figure right).

5.12 DMS, OCS, CS₂ and Isoprene

(D. Booge¹, H. Feil¹, C. Marandino¹, S. Rolfes², B. Engelen²)

¹ GEOMAR, ² Univ. Oldenburg

5.12.1 Introduction

The goal of our project was to understand the impact of anthropogenic inputs on microbial processes and the subsequent effects on the cycling of climate-active trace gases between the ocean and atmosphere. By investigating the aerosols on land and over the ocean (in cooperation with TROPOS), in conjunction with atmospheric trajectories, microbial mediated transformations (in cooperation with WG Engel), trace gas distributions in the ocean and in the atmosphere (in cooperation with WG Quack), we cover the full feedback cycle. The analysis of microbial communities in natural Bay of Bengal water will help to understand the influence of the anthropogenic inputs on the sulphur cycle, specifically the production of the climate-active gas DMS, as well as other biogenically produced and consumed volatile gases such as isoprene, OCS and CS₂, which are measured in the surface ocean (underway) but also in different waters depth during stations (CTD/rosette). Comparing microbial communities from areas with high anthropogenic inputs (northern area of the cruise track) with areas of low inputs (southern area of the cruise track) will give information about the influence of these factors on key organisms involved in the production and degradation of the volatiles. Further, identification of key genes within respective metagenomes will allow us to follow metabolic pathways for the conversion of e.g. organic sulphur compounds. Additionally, we perform incubation experiments on board SO305, in which natural water samples will be spiked with collected atmospheric aerosols from filters or scrubber water from ships. These experiments will allow us to specifically target processes and microorganisms that are involved in trace gas cycling under anthropogenic influences.

5.12.2 Methods

Surface water from the underway pump, located at about 6 m depth in the ship's moonpool, was sampled every 1 to 3 hours, in order to investigate the spatial distribution of DMS, isoprene and CS₂ in the surface waters along the cruise track. Chromophoric dissolved organic matter (CDOM) samples were taken every 3 hours. Besides surface measurements, discrete samples from different depths (surface to bottom) were collected to resolve trace and CDOM concentration variations also in a vertical resolution. The water for the depth profiles were sampled using Niskin bottles attached to the CTD. During each shallow CTD shallow cast, 6 - 7 different depths were sampled. 1 - 2 different depths were sampled from the deep CTD cast. During 4 stations, 5 to 6 different depths were additionally sampled for flowcytometry and DNA/RNA analysis.

Continuous measurements (LGR OCS Analyzer) for OCS/CO₂/CO were carried out in the air (air inlet in front of the bridge) and in the surface water using the ship's underway pump. The one hourly time interval was set to 50 min of water measurements and 10 min of air measurements. Daily integrated aerosol filter measurements were performed from the ship's monkey bridge and were/will be used for incubation experiments on board and back in Germany.

In addition to oceanic and atmospheric samplings, two incubation experiments were performed during SO305. 63 polycarbonate bottles (2 L volume) were filled with surface water from 5 m depth. 21 bottles were spiked with 1% and 2% of scrubber water and with 1% scrubber water and aerosol filter leachate at the beginning of the first and second experiment, respectively. The other 21 bottles were used as controls. The bottles were placed in a water bath at current ocean surface temperature and were exposed to a natural day/night light cycle to a maximum time of six days. Triplicate sample bottles for each treatment (treatment and control) were taken once a day at the same time in order to be analysed for following different parameters: CS₂, isoprene, DMS/DMSP/DMSO, nutrients, DNA, RNA, bacteria and phytoplankton counts, CDOM and trace metals.

All trace gas samples were directly measured on board, within two hours after sampling, using a purge and trap system coupled to a gas chromatograph attached with a mass spectrometer (P&T-GC-MS). After the analysis, samples were prepared to be brought back to GEOMAR home lab for further analysis of DMSP and DMSO concentrations. Besides nutrient measurements, which were performed on board, all other samples will be analysed after the cruise in the home labs of GEOMAR and Univ. Oldenburg.

Originally, a third incubation experiment was planned for the last week of the cruise. Due to the prematurely termination of the cruise we were not able to perform the experiment. Measurements of samples back in the home labs and further data analyses will be finalized within one and a half year after the cruise until end of October 2025.

5.12.3 List of samples/measurements

<u>OCS/CO/CO</u>₂: Continuous measurements; April 17th – May 11th; 50 min water (from UW pump at ~5 m depth), 10 min air (from atmospheric line; inlet above bridge) <u>Aerosols:</u> 24h integrated PM10 aerosol filter sampling; April 17th – May 10th DMS, isoprene and CS₂: Surface water from the underway pump, sampled every 1 to 3 hours Besides surface measurements, discrete samples from different depths (surface to bottom) were collected for trace gases During each shallow CTD shallow cast, 6 - 7 different depths were sampled. 1 - 2 different depths were sampled from the deep CTD cast.

CDOM: samples were taken every 3 hours from the underway pump system.

Table 5.0	Sampling for DIVA/KIVA and noweytoniculy					
Gear	Station	Bedford No.	depth [m]	lat	lon	date
CTD	8_1	30510263	4191.3	3.00	88.00	20.04.2024
CTD	8_1	30510273	759	3.00	88.00	20.04.2024
CTD	8_1	30510284	340	3.00	88.00	20.04.2024
CTD	8_3	30510288	229.1	3.00	88.00	20.04.2024
CTD	8_3	30510297	55.1	3.00	88.00	20.04.2024
CTD	8_3	30510303	2.4	3.00	88.00	20.04.2024
CTD	20_1	30511008	3216.8	12.30	86.00	28.04.2024
CTD	20_4	30511015	1000	12.30	86.00	28.04.2024
CTD	20_4	30511055	240	12.30	86.00	28.04.2024
CTD	20_4	30511058	170	12.30	86.00	28.04.2024
CTD	20_4	30511066	80	12.30	86.00	28.04.2024
CTD	20_4	30511068	1	12.30	86.00	28.04.2024
CTD	25_1	30511254	3167.3	13.00	86.00	01.05.2024
CTD	25_1	30511263	1000	13.00	86.00	01.05.2024
CTD	25_3	30511286	121.4	13.00	86.00	01.05.2024
CTD	25_3	30511289	69.9	13.00	86.00	01.05.2024
CTD	25_3	30511293	1.9	13.00	86.00	01.05.2024
CTD	38_2	30512306	2523.4	16.00	89.60	10.05.2024
CTD	38_4	30512329	200	16.00	89.60	10.05.2024
CTD	38_4	30512335	139.3	16.00	89.60	11.05.2024
CTD	38_4	30512340	36	16.00	89.60	11.05.2024
CTD	38_4	30512346	4.2	16.00	89.60	11.05.2024

 Table 5.8
 Sampling for DNA/RNA and flowcytometry

5.12.4 Preliminary results

First preliminary results show contrasting variabilities of different trace gases in the surface ocean as shown in Figure 5.5. Isoprene, directly produced by phytoplankton, shows slightly elevated concentrations between 6°N and 10°N. Further northwest, values are very low and close to the detection limit indicating very low isoprene production in this area. DMS, a secondary product of microbial production, has highest concentrations at the equator, which could be due to equatorial upwelling of nutrients. Further north, DMS variability follows a similar pattern as isoprene concentrations. CS_2 concentrations are generally low and do not show much variability, except higher concentrations south of the equator. Overall low concentrations of CS_2 indicate low concentrations of CDOM, which is a precursor of photochemical production of CS_2 .

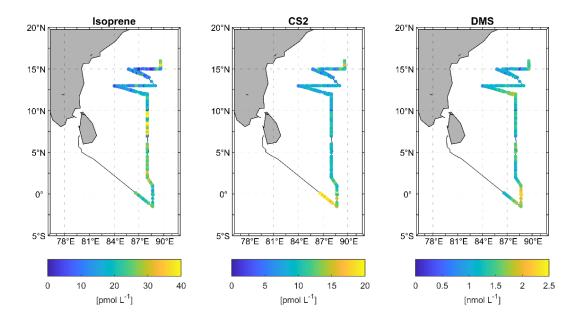


Fig. 5.5 Surface concentrations of isoprene (left), CS₂ (middle) and DMS (right) during SO305.

First insights into the results of one incubation experiment indicate a significant influence of aerosol filter and scrubber water addition on the production of DMS and isoprene (Figure 5.6). Isoprene concentrations decrease over the course of the incubation in the experiment in the control and in the scrubber water treatment. However, after day 3, isoprene concentrations increase in the aerosol filter treatment, which is most likely due to an induced phytoplankton bloom triggered by nitrate addition from the aerosol filters. DMS concentrations show a distinct different picture in the incubation experiment. DMS concentrations increase after day 3 in all three treatments. However, the increase is much lower and higher in the scrubber and aerosol filter treatment, respectively. Increasing concentrations in the aerosol filter treatment are due to increased nitrate concentrations. Lower concentrations in the scrubber water treatment indicate a potential toxic effect of scrubber water constituents on the microbial community and the subsequent biogenic production of DMS.

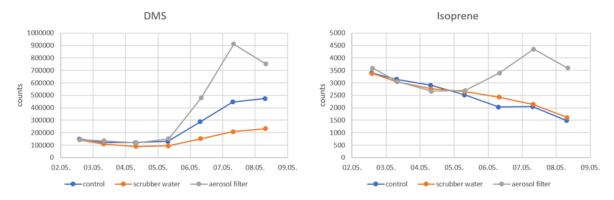


Fig. 5.6 DMS (left) and isoprene (right) values over the course of one experiment using scrubber water and aerosol filter leachate.

5.13 Halogenated Methanes

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5.13.1 Introduction

The greenhouse gas ozone acts as an important UV radiation-protector in the stratosphere for life on earth. It is rapidly destroyed by radical reactions of halogens, e.g. chlorine from long-lived fluorochlorocarbons, which were banned by the Montreal protocol in the 1990s. Ozone is recovering in the global upper stratosphere, while it is still declining in the lower stratosphere of the tropics. Surface ozone, on the other hand, is increasing due to air pollution, especially when fossil fuel is combusted. Over the oceans an active halogen chemistry caused by natural halogenated volatile substances and aerosols destroys ozone effectively. These substances, such as bromoform (CHBr₃), the largest organic bromine source for the atmosphere, dibromomethane (CH₂Br₂) and methyl iodide (CH₃I) are formed in the oceans and contribute to ozone depletion in the troposphere and stratosphere. Despite their short lifetimes, very short-lived substances (VSLS) can be rapidly lifted to the stratosphere by tropical deep convection. Shipboard observations have shown that emissions are often higher near the coast and in upwelling areas than in the open ocean, due to natural sources as phytoplankton and macroalgae. Meanwhile coastal anthropogenic sources must also be taken into account (Mehlmann et al., 2020), as e.g. CHBr₃ is produced in large quantities during the disinfection of seawater and transported to the upper troposphere (Jia et al., 2023). Chlorinated very short-lived substances (VSLS) are identified as present major ozonedepleting VSLS due to increasing anthropogenic emissions (Villamayor et al., 2023). Dichloromethane (CH₂Cl₂) is the most abundant, with the fastest increasing emissions at present, especially from Asia, and emissions of chloroform (CHCl₃) have also grown. Measurements of VSLS in and above the world's oceans are sparse and data show great variability. Little is known about the emission strengths or the uptake into the seawater and biogeochemical cycling of VSLS from the Indian Ocean. A single measurement campaign in the Bay of Bengal (Yamamoto et al., 2001) suggests that the Bay of Bengal might be a significant source of bromine and iodine to the atmosphere, as concentrations of the compounds were generally higher than data reported for other open ocean regions at mid- and high-latitudes (Ziska et al., 2013). During SO305, we measured the above compounds in water and air, in order to understand their source strengths and learn about their biogeochemical cycling in the oxygen minimum zone of the Bay of Bengal.

5.13.2 Methods

Sampling of the halocarbons in seawater was done with 250 ml amber glass bottles from the CTDrosette bottles and from the underway supply by the submersible pump in six to twelve-hour intervals, which were filled bubble free and closed with Teflon lids. Surface water samples were stored in the lab, deep water samples in a \sim 7°C refrigerator until they were measured with a purge and trap system attached to a gas chromatograph (Agilent 6890N) with mass spectrometric detection (Agilent Technologies 5975 inert XL MSD) (GC-MS), within 1-12 hours after the sampling. Prior to analysis, the samples were warmed to room temperature (\sim 20°C) to ensure purge efficiency and consistency. The bottles were attached to a manifold, which pressurizes the

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top of the sample with helium and fills the purge-chamber with 30 ml of the seawater through a second submerged outlet, so that a pure sample can by transferred. Halocarbon concentrations were measured by purging 38 ml of sample water from ambient to 65°C in a long glass chamber for 45 min with a stream of helium at around 52 ml min-1, resulting in a purge efficiency of > 98%. A Nafion dryer (Ansyco GmbH, TT-050) was installed for the desiccation of the purge flow, using a counter-flow of 120 ml min-1 of N2. The volatilized trace gases were trapped between -150 and -170 °C above liquid nitrogen in a coil of deactivated stainless-steel tubing. After purging, the compounds were desorbed from the trap and injected into the GC-MS by heating the trap electronically to 180°C within 1.5 min. The sample was then separated on a 60 m by 0.38 mm Rtx-624 capillary column (film thickness = $1.8 \,\mu$ m) and finally detected by the MS in single ion mode. The concentrations were quantified with external standards, which were volumetrically prepared in methanol. The calibration curves were prepared with 1-4 µl of standards containing 0.02, 0.3 and 3 pmol of compound per µl. During transport of the standards, contamination occurred within the standard flasks and they will be recalibrated in Kiel. One calibration standard was repeatedly run in triplicate for sample quantitation, to account for sensitivity changes of the mass spectrometer during the cruise. The seawater measurements generally have a reproducibility of 2 to 10% for methyl iodide (CH₃I), dichloromethane (CH₂Cl₂), chloroform (CHCl₃), carbon tetrachloride (CCl₄), dibromomethane (CH₂Br₂), chloroiodomethane (CH₂ClI), trichloroethene (CHCl₃), tetrachloroethene (C₂Cl₄), dichlorobromomethane (CHCl₂Br), dibromochloromethane (CHBr₂Cl), bromoform (CHBr₃), diiodomethane (CH₂I₂).

Underway air samples of SO305 were taken at the bow in 2 L stainless steel canisters with a metal bellows pump. Air samples will be analysed for halocarbons at the National Center for Atmospheric Research (NCAR) in Boulder by Elliot Atlas (RSMAS, Miami) using GC/MS. The samples were taken simultaneously with underway seawater samples or with a delay of up to two hours to the corresponding water samples. Air samples were taken in parallel to the NCAR-samples in order to intercalibrate between air and water measurements and those samples will be analysed in Kiel on the onboard GC/MS.

Air-sea fluxes of the compounds will be calculated using air-sea exchange parameterizations. Compound specific transfer coefficients (k_w) will be derived from the transfer coefficient k_{CO2} of CO₂ by Schmidt number corrections (Sc).

All data will be available in winter 2024/2025.

5.13.3 Phytoplankton Pigments

Water samples taken from 6 surface depth (0-200 m) of the CTD deployments and regularly from 6h intervals from the underway sampling have been filtered for phytoplankton pigment analysis by HPLC for later biomass and algal taxonomic group analysis. Flow cytometry subsamples (850 μ L) have been taken from the individual water samples and where added to Glutaraldehyde in 1 mL vials. After 30 minutes at 4°C the samples were frozen in liquid nitrogen and then stored at -80°C for later analysis in the Biological Oceanography at GEOMAR. 2 L of water samples were immediately filtered through GF/F-Filters, at an under pressure of maximum 200 mbar and the filters stored at -80°C until later analysis in the Biological Oceanography at GEOMAR.

5.13.4 List of Samples

382 seawater samples from depth profiles (see Table 11.3 in the Appendix) and 89 surface seawater samples were taken (see Table 11.4 in the Appendix) and analysed on board. 80 air sample canisters for Elliot Atlas (RSMAS, Miami, FL, USA) to be measured at NCAR, Boulder and 20 air samples for intercalibration in Kiel were taken during the cruise (see Table 11.5 in the Appendix). A list of samples for phytoplankton pigment analysis is given in Table 11.6. in the Appendix.

5.13.5 Preliminary Results

Varying distributions of the 12 halocarbons in surface and deep waters of the Bay of Bengal, reveal patterns of the compounds, related to biology, water masses and air-sea gas exchange. The following plots (Fig. 5.7 - 5.10) show some results of a few halocarbons in the upper 600 m of the water column.

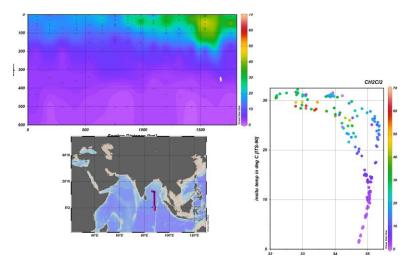


Fig. 5.7 Dichloromethane (CH₂Cl₂) is an anthropogenic compound, with increasing concentrations in the atmosphere. It increases in less saltier waters towards the northern Bay of Bengal.

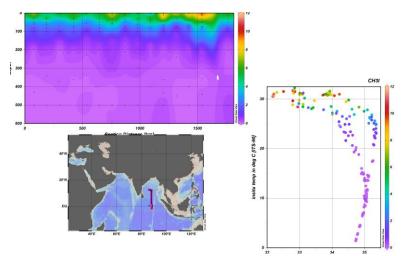


Fig. 5.8 Methyl iodide (CH₃I) is produced in sunlit reactions with DOM. Its concentrations are higher during the day in surface waters and appear to also increase in less saline surface waters.

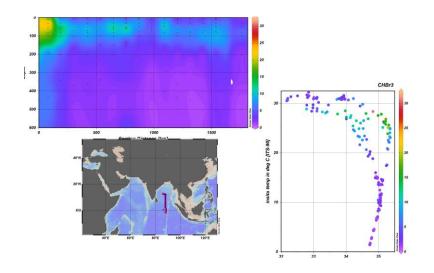


Fig. 5.9 Bromoform (CHBr₃) is produced in the chlorophyll maximum of the entire oceans. It appears to have higher concentrations in the equatorial region.

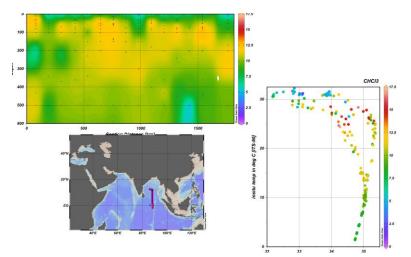


Fig. 5.10 Chloroform (CHCl₃) is both a natural and anthropogenic compound. It appeared to be elevated in a certain water mass, below the mixed layer.

5.14 Nitrogen Cycle - Isotopes

(G. Schulz¹, L. Nazzari¹, T. Sanders¹, K. Dähnke², N. Lahajnar¹, B. Gaye¹) ¹ Univ. Hamburg, ² Hereon

5.14.1 Introduction

The SO305 BIOCAT-IIOE2 project aims to decipherer nitrogen sources as well as turnover processes to contribute to a better understanding of the emergence and spreading of the oxygen minimum zone in the Bay of Bengal. The working groups of Biogeochemistry in the Earth System from the University of Hamburg and Aquatic Nutrient Cycles from the Helmholtz Centre Hereon contribute to this goal using a two-fold approach by (1) sampling natural abundance isotopes of reactive nitrogen in the water column and (2) conducting process rate studies for nitrification and nitrogen and carbon uptake.

Accurate rate measurements are rare in oxygen minimum zones, and the understanding of the oxygen threshold values of different nitrogen turnover processes is constantly evolving. This also

applies to the understanding of nitrification, which can occur in niches with very low oxygen concentrations, partially inhibited by oxygen, and is therefore not realistically reproduced in models (Sun et al., 2021). Changes in nitrate stable isotopes have been used frequently to track sources and sinks of reactive nitrogen (e.g. nitrification, N₂ fixation, assimilation, and denitrification) in the open ocean (e.g. Sigman et al., 2005; Gaye et al., 2013; Marconi et al., 2015; Harms et al., 2019). Biological turnover processes favour lighter isotopes, leading to an enrichment in the remaining nitrogen pool (Granger et al., 2004; Dähnke et al., 2010; Sigman and Fripat, 2019). Thus, elevated nitrate isotope ratios in the photic upper layers are a possible sign for assimilation and can indicate consumption via denitrification and/or anammox in the oxygen minimum zone (e.g. Granger et al., 2004; Sigman and Fripat, 2019).

5.14.2 Sampling Strategy

For natural abundance isotopes, water samples were filtered (CN/GF, MINISART, 0.45 μ m) and stored frozen for later nitrate analysis in the home lab. For nitrite isotope analysis, 1000 μ L of 1M KOH was added to one sample replicate and kept cool to minimize oxygen isotope exchange (Böhlke et al., 2007). To quantify nitrification, we used a parallel setup of the ammonium tracer method (Ward, 2011) and the isotope dilution method (Norton and Stark, 2011; Sanders et al., 2018). Both methods are based on nitrate stable isotope measurements, whereas the ammonium tracer method tracks the turnover of ¹⁵N-labelled ammonium to nitrate and the dilution method the isotopic dilution of a ¹⁵N-labelled nitrate pool by freshly formed ¹⁴N-nitrate. Incubation time was 24 hours and up to 10 days for the ammonium method and isotope dilution method, respectively. Subsamples at different time steps were filtered (CN/GF, MINISART, 0.45 μ m) and stored frozen for later nutrient and nitrate isotope analysis.

Water samples for uptake incubations were spiked with NaH¹³CO₃ and either ¹⁵N-nitrate or ¹⁵N-ammonium. After an incubation time of 2 and 4 hours for ammonium and nitrate uptake, respectively, at in situ light and temperature conditions, triplicate water samples were filtered over pre-combusted GF/F filters (450 °C, overnight). Filters were dried (40 °C, 48 hours) and stored in the dark for later analysis of ¹⁵N and ¹³C of the suspended particulate matter. Samples will be measured and data finalized in the following two years.

5.14.3 List of Samples

In total, natural abundance stable isotope samples were taken at 21 stations (Tab. 5.9) from 39 CTD casts with a high resolution of 25 - 50 m in water depths with low oxygen concentrations. Incubation experiments to determine nitrification rates in the water column were done at 6 stations at five to six distinct water depths chosen based on oxygen and fluorescence profiles. Nitrogen and carbon uptake experiments were done at 5 stations in two water depths.

Gear	Station	Date	Latitude	Longitude		Samples	
					NA	NP	UR
CTD	2	17.04.2024	0.6°S	88.4°E	25		
CTD	4	18.04.2024	0.0°S	88.4°E	27	5	
CTD	6	19.04.2024	1.0°N	88.4°E	22		
CTD	8	20.04.2024	2.6°N	87.6°E	23		
CTD	10	22.04.2024	4.6°N	87.6°E	24	6	2
CTD	11	22.04.2024	5.6°N	88.0°E	23		
CTD	15	25.04.2024	9.0°N	88.0°E	21		
CTD	18	26.04.2024	12.0°N	88.0°E	23	6	2
CTD	20	28.04.2024	12.3°N	85.6°E	22		
CTD	23	30.04.2024	13.0°N	84.0°E	25		
CTD	24	30.04.2024	13.0°N	85.0°E	6		
CTD	25	01.05.2024	13.0°N	86.0°E	9		
CTD	26	01.05.2024	13.0°N	87.0°E	6		
CTD	27	02.05.2024	12.6°N	88.0°E	24		
CTD	28	03.05.2024	12.6°N	89.0°E	6		
CTD	30	04.05.2024	13.6°N	88.0°E	24	6	2
CTD	32	06.05.2024	14.6°N	85.4°E	26	6	2
CTD	34	08.05.2024	15.0°N	86.6°E	25		
CTD	35	08.05.2024	15.0°N	87.6°E	6		
CTD	37	09.05.2024	15.0°N	89.4°E	25	6	2
CTD	38	10.05.2024	15.6°N	89.4°E	24		

Table 5.9 Overview of natural abundance isotopes (NA), nitrification potential (NP) and uptake rates (UR) sampling depths.

5.14.4 Preliminary Results

The isotopic composition of natural abundance samples and process rates results will only be available after analysis in the home lab. However, nutrient data shows accumulation of nitrate concentration in most sampled depths, which will allow isotopic determination. The dual stable isotope approach comparing $\delta^{15}N$ and $\delta^{18}O$ of nitrate will help to identify nitrate assimilation in the photic upper layer as well as nitrate consumption (denitrification and/or anammox) in the regions with low oxygen concentrations (e.g. Granger et al., 2004; Sigman and Fripat, 2019). Within the primary oxygen minimum zone, a portion of the samples exhibited nitrite accumulation. Conducting separate nitrite isotope analysis will help in disentangle the processes accountable for nitrite accumulation in these samples. Additional interpretation and process rate calculations will be feasible following isotopic determination.

5.15 Suspended Matter

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¹ Univ. Hamburg

5.15.1 Introduction

The objective of the SO305 BIOCAT-IIOE2 project is to investigate the sources and degradability of particulate organic matter, with the goal of enhancing our comprehension of the development and expansion of the oxygen minimum zone in the Bay of Bengal. The working group Biogeochemistry in the Earth System from the University of Hamburg contribute to the project by collecting suspended matter samples.

Suspended particulate matter samples help to assess the composition of non-sinking particulate matter and advance our understanding of biogeochemical element cycles. Since suspended matter is too small and insufficiently dense to sink, it primarily disperses into deeper waters through the subduction of surface waters (Resplandy et al., 2019) and is passively transported by the ocean conveyor belt (Silver et al., 1998; McCave, 1984). Consequently, suspended matter sampling allows to trace the biogeochemical processes of source water masses and thereby decipher water mass transport mechanisms in the research area.

5.15.2 Methods

Suspended particulate matter was sampled via large-volume filtration of water samples collected at selected depths using the CTD/rosette with 24 10 L bottles. Water samples were filtered using a self-made filtration unit (Universität Hamburg) and combusted (450 °C, overnight), pre-weighed GF/F filters (WHATMAN, ~0.7 μ m, 47 mm diameter). Following filtration, the suspended matter samples were rinsed with MILLI-Q water to remove sea salt. Afterwards, filters were dried at 40 °C for 48 hours in the ship's dry oven and stored in a dark. In the home lab, suspended particulate matter composition will be analysed starting with suspended particulate matter load followed by analyses of total nitrogen, total carbon, organic carbon, amino acid composition and stable isotope ratios of nitrogen. Samples will be measured and data finalized in the following two years.

5.15.3 List of Samples

In total, suspended particulate matter samples were taken at 23 stations leading to 165 filters (Table 5.10). Samples were taken at distinct depths depending on the CTD profile ranging between five and nine depths per station.

Gear	Station	Date	Latitude	Longitude	Depths
	2				-
CTD		17.04.2024	0.6°S	88.4°E	5
CTD	4	18.04.2024	0.0°S	88.4°E	5
CTD	6	19.04.2024	1.0°N	88.4°E	5
CTD	8	20.04.2024	2.6°N	87.6°E	5
CTD	10	22.04.2024	4.6°N	87.6°E	5
CTD	11	22.04.2024	5.6°N	88.0°E	5
CTD	13	24.04.2024	7.0°N	88.0°E	5
CTD	15	25.04.2024	9.0°N	88.0°E	5
CTD	18	26.04.2024	12.0°N	88.0°E	8
CTD	20	28.04.2024	12.3°N	85.6°E	5
CTD	23	30.04.2024	13.0°N	84.0°E	9
CTD	24	30.04.2024	13.0°N	85.0°E	7
CTD	25	01.05.2024	13.0°N	86.0°E	9
CTD	26	01.05.2024	13.0°N	87.0°E	7
CTD	27	02.05.2024	12.6°N	88.0°E	9
CTD	28	03.05.2024	12.6°N	89.0°E	7
CTD	30	04.05.2024	13.6°N	88.0°E	9
CTD	32	06.05.2024	14.6°N	85.4°E	8
CTD	34	08.05.2024	15.0°N	86.6°E	6
CTD	35	08.05.2024	15.0°N	87.6°E	7
CTD	36	09.05.2024	14.6°N	88.4°E	6
CTD	37	09.05.2024	15.0°N	89.4°E	9
CTD	38	10.05.2024	15.6°N	89.4°E	4

Table 5.10Overview of suspended matter sampling.

5.15.4 Preliminary Results

The composition of the suspended particulate matter will only be available after analysis in the home lab. Visual inspection of loaded filters showed generally higher loaded filters in the shallower water depths. The filters from the fluorescence maximum had the highest loads with a greenish colour.

5.16 Nitrogen Cycle - Microbial Processes

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5.16.1 Introduction

During the SO305 BIOCAT-IIOE2 cruise, water was collected for on board incubations using various ¹⁵N-nitrogen species combined with water filtrations for posterior molecular analysis in order to elucidate potential rates of microbial nitrogen transformation processes, and the key players involved in such processes. While other oxygen minimum zones (OMZ) in regions such

as the East Pacific and the Arabian Sea experience an annual estimated net loss of nitrogen to the atmosphere (14 Tg N yr⁻¹), the Bay of Bengal (BoB) exhibits a significantly lower net loss (7 Tg N yr⁻¹) compared to other systems. This lower loss can be attributed to the unique characteristics of the Bay of Bengal, which may account for the trace amounts of oxygen found in the area. Therefore, our study aims to gain a deeper understanding of the processes controlling nitrogen cycling in the BoB.

5.16.2 Methods

On board the RV SONNE, incubations were carried out in deck-based tanks as well as in the climate lab using stable isotope tracing. The deck-based incubations contained water sampled at the surface, chl-max and O_2 gradient. The sampled water was amended with ${}^{15}N_2$ gas to quantify the potential rate of nitrogen fixation over a period of 24 hours in addition to ${}^{13}C$ -bicarbonate to determine the potential rate of carbon fixation.

Concurrently, experiments were carried out in the temperature-controlled lab using ¹⁵Nnitrogen species quantifying processes responsible for the loss of nitrogen through denitrification (¹⁵NO₂⁻, ¹⁵NO₃⁻, and ¹⁵N₂O) and anammox (¹⁵NH₄⁺ and ¹⁵NO₂⁻) or cycling of nitrogen through oxidation processes (¹⁵NH₄⁺ and ¹⁵NO₂⁻). Kinetic experiments using either ¹⁵N-nitrogen species or oxygen were also carried to determine the role these environmental factors for microbial nitrogen cycling in the BoB.

Additional water was collected from incubation depths and filtered for DNA and RNA extraction and will be used for metagenomic and transcriptomic analysis. In addition to this, water was also collected and filtered for Fluorescence in Situ Hybridization (FISH) analysis. The collected samples will be processed at SDU using techniques such as GC-IRMS and elemental analyser/MS for N-cycling experiments. Molecular work will involve FISH, DNA and RNA sequencing, as well as flow cytometry.

Unfortunately, the temperature lab did not work optimally with temperature fluctuating between 8°C to 18°C (target temperature 12°C). This was a problem for the in-lab incubations. It will take approx. 1 year to get all data from the experiments conducted in the climate lab.

5.16.3 List of Samples

Water was collected from the CTD and the underway pump. All 24h-stations were targeted for incubations, while additional stations were sampled occasionally for both incubation work and molecular samples. Molecular samples were collected across the cruise track, while incubation experiments started from 5°N (station 10). The primary focus for the incubation experiments were on the northern most stations where the OMZ was located. In each CTD, samples were collected from surface, chlorophyll max, and several depths between 100 and 400 covering the OMZ for incubation and molecular analyses.

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Station	Molecular	N ₂ -fixation	N-cycling	Date
4	X			18/04/2024
6	X			19/04/2024
8	X			20/04/2024
10	X	X	Х	21/04/2024
13	X	X	Х	24/04/2024
15	Х			25/04/2024
16	Х		Х	26/04/2024
18	X	X	Х	27/04/2024
20			Х	28/04/2024
22	Х			29/04/2024
23	Х	Х	Х	30/04/2024
26	х		Х	02/04/2024
28	x			03/04/2024
30	X	X	X	04/04/2024
32	X	X	Х	07/04/2024
34	X			08/04/2024
35	X		X	09/04/2024
37	X	Х	X	10/04/2024

Table 5.11- Stations sampled for N- and C-fixation (on-deck experiments) and molecular work.

Stations sampled for N cycling and N loss (temperature lab experiments): 10, 13, 16, 18, 20, 23, 26, 30, 32, 35, and 37. Number of samples generated: For the recycling vs. loss of fix nitrogen analyses we generated a total of 876 samples. For FISH samples, we generated a total of 72 samples. For N_2O analysis we generated xxx samples. For N_2 fixation we generated a total of xxx samples.

5.16.4 Preliminary Results

Since all the samples collected have to be processed at SDU, we cannot not show any preliminary result.

5.17 Drifting Sediment Traps

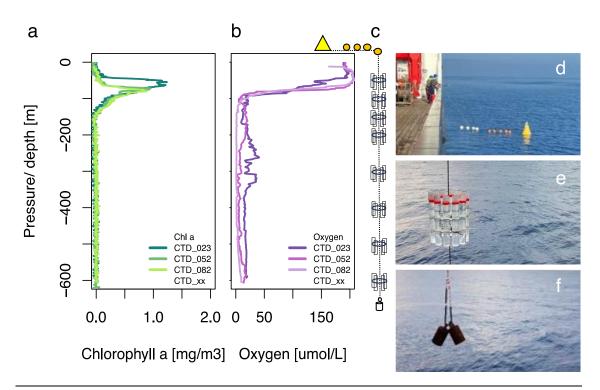
(T. Barthelmeβ¹, J. Roa¹, S. Golde¹, A. Barbot¹, B. Pontiller¹, K. Becker¹, A. Engel¹)
¹ GEOMAR

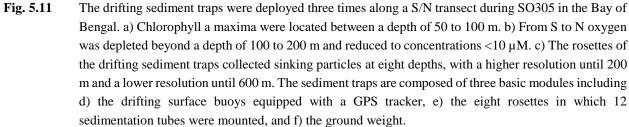
5.17.1 Introduction

The Bay of Bengal in the Indian Ocean is an oligotroph system, which is defined by complex eddies, currents, and divergent water mass stratification in which a deep Chlorophyll a maximum evolves between 50 and 100 m depth (Fig. 5.11a). At the northern stations of the cruise track of SO305, the surface waters are expected to be influenced by the enormous freshwater inflow of the Brahmaputra and Ganges delta. The Bay of Bengal is furthermore characterized by an expanded oxygen minimum zone (OMZ) below a varying depth of 100 to 200 m (Fig. 5.11b). Depending on

the stations, it expands towards more than 600 m depth. OMZs are thought to evolve due to the degradation of organic matter via respiration of organisms, i.e. the consumption of oxygen, and poor physical ventilation. Although primary production in the surface waters was shown to be rather low during the last decades, the export of organic carbon towards depth is larger than expected in the Bay of Bengal (Unger and Jennerjahn, 2009; Rixen et al., 2009). Hypothetically, riverine organic matter and inflow of lithogenic material may 1) enhance particle formation, and 2) accelerate particle sedimentation by ballasting (Ray et al., 2015). Additionally, 3) polluted air masses have been recently shown to form a large cloud which may transport minerals, nutrients, and pollutants off the coast (Ramanathan et al., 2017). Pollutants may deposit at the ocean surface and are considered as a further factor influencing surface biogeochemistry and particle export. In general, data are scarce which assess particle export and export efficiency. They are yet insufficient to fully comprehend the evolution of OMZ and their influence on long-term carbon storage (Engel et al., 2023).

To assess biogenic particle export dynamics out of the surface mixed layer into the OMZ, we deployed drifting sediment traps along a S/N transect at 88°E from 5°N to 15°N. The drifting sediment traps resolve the sedimentation flux at high resolution in the surface ocean (every 50 m) until a depth of 200 m and at lower resolution (every 100 m) down to a depth of 600 m (Fig. 5.11c). The drifting buoys at the surface hold the sediment traps in place which are further ballasted by a ground weight (Fig. 5.11d-f). The most important components of the traps are the eight rosettes equipped with twelve sedimentation tubes. Above a formaldehyde-brine at the bottom, the tubes are filled with filtered seawater (Fig. 5.11e). The brine conserves the captured, sinking particles. The sediment traps deployments lasted between 48 and 69 hours. To define particle export dynamics, we have collected samples for the analysis of e.g., particulate organic carbon (POC) and its molecular components including lipids, amino acids, and carbohydrates. Samples for biogenic silica (BSi) and total particle mass were also collected, enabling us to deduce the contribution of lithogenic material. The contribution of different organisms to particle formation and degradation along its sedimentation pathway is complemented by enzyme activity measurements, meta-genome, and -transcriptome analysis. To assess the surrounding water column and its potential POC budget driving sedimentation fluxes, additional CTD casts were sampled at the beginning and end of each sediment trap deployment and at in total nine different depths (additionally including 5 m depth sample).





5.17.2 Methods

We collected mainly particulate organic matter (POM) larger than 0.7 µm in size, exceptions are mentioned in the paragraph below. The material collected in the sedimentation tubes was pooled, well-mixed and distributed for triplicate filtration. Samples for POC, particulate organic nitrogen (POC), total particulate carbon (TPC), particulate amino acids (PAA), particulate carbohydrates (PCHO), Chlorophyll a (Chl a), and particulate organic phosphorus (POP) were filtered onto GFF filters (Whatman, 0.7 µm), while samples for particulate lipids were filtered onto PVDF membranes (Millipore, 0.2 µm), biogenic silica (BSi) on cellulose acetate filters (Sartorius, 0.8 µm), and total particle mass, transparent exopolymer particles (TEP), and Coomassie stainable particles (CSP) were filtered onto polycarbonate membranes (Whatman, 0.45 µm pore size) applying a vacuum of ~200 mbar. Leftover sample volume was filled into glass bottles for the analysis of microplastics. In order to asses meta-transcriptomics (meta-t, based on RNA) and genomics (meta-g, based on DNA), two size fractions were distinguished and filtered onto Sterivex (0.22 µm) and PVDF membranes (Millipore, 3 µm) for selected depths (50, 300 and 600 m). Extracellular enzyme activity (EEA) was assessed for the same selected depths from CTD casts and trap deployments. Selected sedimentation tubes were further prepped to collect sinking particles in a gel-matrix for microscopical analysis (conserved in polyacrylamide, Lundsgaard,

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1995; Engel et al., 2022). Flow cytometry samples were collected for the analysis of phytoplankton cell abundance, bacterial cell counts, and virus particles in addition to POM from corresponding CTD casts.

In brief, POC, PON, and TPC is analysed using a Euro EA elemental analyser (Sharp, 1974). PAA and PCHO will be determined by high-performance liquid chromatography (HPLC) and anion exchange chromatography (HPAEC), equipped with a fluorescence and pulsed amperometric detector (PAD), respectively (Lindroth and Mopper, 1979; Engel and Händel, 2011). In total, 13 amino acids and 12 carbohydrate monomers can be discriminated including neutral and acidic compounds. Lipids and pigments will be analysed on a HPLC coupled to a Orbitrap high resolution mass spectrometer (MS) according to Becker et al. (2018). Chl a will be assessed via photometry (Turner Design, USA) after the modified protocol of Evans et al. (1987). After digestion and dissolution, POP and BSi will be analysed in our central nutrient lab using photometry (QuAAtro, Seal Analytical; Grasshoff et al., 1999). Total particle mass will be assessed by high-precision weight. TEP and CSP were stained with Alcian and Coomassie Blue after filtration, which enables the distinction of carbohydrate-like and protein-like particles via photometry (Passow and Alldredge, 1995; Cisternas-Nova et al., 2014). Microplastics will be analysed by digestion (Cole et al., 2014; Mintening et al., 2016), followed by pyrolysis and coupled to gas chromatography and MS (GC-MS) (Fischer et al., 2019). Meta-g and meta-t analysis will follow the methodology as outlined by Fontanez et al., (2015). EEA produced by the overall microbial community was assessed on board by spectrofluorometric measurements of fluorogenic substrates. Following the methodology outlined by Hoppe (1983), the hydrolysis rates of five distinct enzymes were determined, including e.g., peptidase and glucosidase activity (Baltar et al., 2010). Flow cytometer samples are fixed in glutaraldehyde and flow cytometry (FACSCalibur) will allow the distinction of five different phytoplankton classes via their cell size distribution (0.2-20 µm) and phytopigments (phycoerythrin and Chl a). To count bacteria and virus particles, samples will be stained with SYBR Green (following standard procedure of our lab, e.g., Engel and Galgani, 2016).

We applied a new sediment trap design during SO305 BIOCAT-IIOE2. After the first deployment (23.04.2024), we lost in total nine out of 96 tubes. During the process of recovery, the sediment traps started spinning due to tension and internal coiling of the rope. The new rosette holders of the tubes could not withstand the centrifugal forces which unravelled. In order to avoid further losses, we implemented three measures, i.e. cutting the rope and integrating vertebra to avoid the internal progression of spin, fixing the tubes to a further rosette holder, and tightening them with cable ties instead of bungee ropes. Our measures were successful.

We also installed an optical fluorescent dissolved organic matter (FDOM) sensor at the CTD frame, however, had to change its orientation from horizontal to vertical (pointing towards depth) as we noticed that scattering of the data was significantly reduced (29.03.2024, 10:43 UTC).

All data measurements will be finalized within two years and selected data are made available on request to interested colleagues. We highly welcome potential collaborations and common publications. After publication in a scientific journal, data will be published in an open-access online repository such as PANGAEA.

5.17.3 List of Samples

The list of samples from the drifting sediment trap deployments is given in Table 11.7 in the Appendix.

5.18 Microbial Activity and Community Composition

(A. Barbot¹, A. Engel¹)

¹ GEOMAR

5.18.1 Methods

The activity of extracellular enzymes (EEA) produced by the overall microbial community was assessed on board by spectrofluorometric measurements of fluorogenic substrates. Following the methodology outlined by Hoppe (1983), the hydrolysis rates of five distinct enzymes were determined, as detailed in the studies conducted by Baltar et al. (2010). To assess the microbial community composition and diversity, filtrations was conducted on two size fraction filters, 0.22 μ m and 0.3 μ m. Metagenomic and meta-transcriptomic analysis will be done within one year following the methodology outlined by Fontanez et al. (2015).

Potential hydrolytic rates of L-Leucin-7-amido-4-methylcoumarin-hydrochlorid (LAPase), 4-Methylumbelliferyl-α-D-glucopyranosid (AGase), 4-Methylumbelliferyl-β-D-glucopyranosid 4-Methylumbelliferyl-N-acetyl-β-D-glucosaminid (NAG), 4-Methylumbelliferyl-(BGase), phosphat (APase) were determined with fluorescent substrate analogs (Hoppe, 1983). 10 µL of the following standards, L-leucine-7-amido-4-methylcoumarin (Sigma Aldrich) and 4methylumbelliferone (Sigma Aldrich) were added with final concentrations of 1.25, 2.5, 10, 50, 100, 500 and 1000 nmol L⁻¹ in black 96 well plates (Costar) and kept frozen for at most one day until replicates of 290 µL water sample were added. These concentrations were previously determined as saturating substrate concentrations (Baltar et al, 2010). The final substrate concentration of 30 µmol L⁻¹ was used to measure Agase, BGase and NAG activities, 100 µmol L^{-1} for APase, and 500 µmol L^{-1} for LAPase. Individual depths corresponding plats were incubated in the dark, close to in situ temperature. After 0, 3, 6 and 12 h of incubation, fluorescence was measured with a plate reader fluorometer (FLUOstar Optima, BMG Labtech, excitation: 355 nm; emission: 460 nm).

Enzyme activities were measured onboard immediately after sampling using L-leucine-4methylcoumarinyl-7-amide (MCA) hydrochloride and 4-methylumbelliferone (MUF) β -Dglucopyranoside (Sigma-Aldrich) as substrate proxies for Lpase (hereafter referred to as peptidase), AGase and BGase activities (hereafter referred to as glucosidase), respectively (Hoppe, 1983). Enzymatic hydrolysis of MCA- and MUF-substrate proxies can be measured with shortterm (several hour) incubations and is generally considered to reflect activities of the in situ microbial community. Fluorescence changes were calibrated using MUF and MCA standard solutions in seawater and used to calculate hydrolysis rates.

5.18.3 List of Samples

Table 5.12	Extracellular ei	nzymatic activi	ty and metage	nomic samples.

Date (UTC)	Time [UTC]	Lat (N)	Long(W)	Station	CTD/TRAPcast	Niskin bottle#1	Niskin bottle#2	depth_m	Comments
23/04/2024	8:06:00 AM	5° 03.344 N	88° 11.796 E	SO305/12-1	TRAP_1			50	Only 0.22 µm filt
23/04/2024	8:36:00 AM	5° 03.344 N	88° 11.796 E	SO305/12-1	TRAP_1			300	
23/04/2024	8:47:00 AM	5° 03.344 N	88° 11.796 E	SO305/12-1	TRAP_1			600	
29/04/2024	1:58:24 AM	11° 56.548 N	87° 38.265 E	SO305/21-1	TRAP_2			50	
29/04/2024	2:25:52 AM	11° 56.548 N	87° 38.265 E	SO305/21-1	TRAP_2			300	
29/04/2024	2:49:23 AM	11° 56.548 N	87° 38.265 E	SO305/21-1	TRAP_2			600	
07/05/2024	11:28:09 AM	13° 58,912' N	088° 13,463' E	SO305/33-1	TRAP_3			50	
07/05/2024	12:03:44 PM	13° 59,019' N	088° 13,649' E	SO305/33-1	TRAP_3			300	
07/05/2024	12:32:52 PM	13° 59,241' N	088° 13,741' E	SO305/33-1	TRAP_3			600	
23/04/2024	8:06:00 AM	5° 03.344 N	88° 11.796 E	SO305/12-1	TRAP_1			50	
23/04/2024		5° 03.344 N	88° 11.796 E	SO305/12-1	TRAP_1			300	
23/04/2024	8:47:00 AM	5° 03.344 N	88° 11.796 E	SO305/12-1	TRAP_1			600	
29/04/2024	1:58:24 AM	11° 56.548 N	87° 38.265 E	SO305/21-1	TRAP_2			50	
29/04/2024	2:25:52 AM	11° 56.548 N	87° 38.265 E	SO305/21-1	TRAP_2			300	
29/04/2024	2:49:23 AM	11° 56.548 N	87° 38.265 E	SO305/21-1	TRAP_2			600	
07/05/2024	11:28:09 AM	13° 58,912' N	088° 13,463' E	SO305/33-1	TRAP_3			50	
07/05/2024	12:03:44 PM	13° 59,019' N		SO305/33-1	TRAP_3			300	
07/05/2024	12:32:52 PM	13° 59,241' N	088° 13,741' E	SO305/33-1	TRAP_3			600	
22/04/2024	5:10:00 AM	4° 59.926 N	87° 59.774 E	SO305/10-13	CTD 10-13	SO305_1_455		50	Deployment C
22/04/2024	5:10:00 AM	4° 59.926 N	87° 59.774 E	SO305/10-13	CTD 10-13	SO305_1_444		300	
22/04/2024	5:10:00 AM	4° 59.926 N	87° 59.774 E	SO305/10-13	CTD 10-13	SO305_1_338	SO305_1_339	600	
27/04/2024	7:40:00 AM		87° 59.933E	SO305/18-8	CTD 18-8	SO305_1_851		50	
27/04/2024	7:40:00 AM	12°00.139 N	87° 59.933E	SO305/18-8	CTD 18-8	SO305_1_842		300	
27/04/2024	7:40:00 AM		87° 59.933E	SO305/18-8	CTD 18-8	SO305_1_834	SO305_1_835	600	
05/05/2024	12:05:00 AM	13° 59.653 N	87° 59.119 E	90305/30-12	CTD 30-12	SO305_1_1709		50	
05/05/2024	12:05:00 AM	13° 59.653 N	87° 59.119 E	90305/30-12	CTD 30-12	SO305_1_1710		300	
05/05/2024	12:05:00 AM	13° 59.653 N	87° 59.119 E	SO305/30-12	CTD 30-12	SO305_1_1092	SO305_1_1093	600	
23/04/2024	9:31:00 AM	5° 03.345 N	88° 11.797 E	SO305/12-2	CTD 12-2	SO305_1_543		50	Recovery C
23/04/2024	9:31:00 AM	5° 03.345 N	88° 11.797 E	SO305/12-2	CTD 12-2	SO305_1_534		300	
23/04/2024	9:31:00 AM	5° 03.345 N	88° 11.797 E	SO305/12-2	CTD 12-2	SO305_1_526	SO305_1_527	600	
29/04/2024	4:00:00 AM		87° 37.442 E	SO305/21-2	CTD 21-2	SO305_1_1093		50	
29/04/2024	4:00:00 AM		87° 37.442 E		CTD 21-2			300	
29/04/2024	4:00:00 AM	11° 56.730 N	87° 37.442 E	SO305/21-2	CTD 21-2			600	
05/05/2024	12:44:00 AM		87° 59,129' E		CTD32-2			50	
05/05/2024	12:44:00 AM	13° 59,650' N	87° 59,129' E	SO305_33_2	CTD32-2			300	
05/05/2024	12:44:00 AM	13° 59,650' N	87° 59,129' E	SO305_33_2	CTD32-2	SO305_1_1934	SO305_1_1935	600	
27/04/2024	2:42:00 AM	12° 00,137' N	087° 59,938' E	90305_18_6	Underway				
10/05/2024	2:14:00 AM	15° 00,140' N	089° 36,124' E	SO305_37_6	Underway				
27/04/2024	2:42:00 AM	12° 00,137' N	087° 59,938' E	SO305_18_6	Zodiac				
10/05/2024	2:14:00 AM	15° 00,140' N	089° 36,124' E	SO305_37_6	Zodiac				
01/05/2024	12:07:00 AM	13° 00,098' N	085° 00,065' E	SO305/24 3	CTD 24-3	SO305_1_1243		50	13° N West -
			085° 00,065' E			SO305_1_1226		300	Trans
			085° 00,065' E	SO305/24_1	CTD 24-1			600	
01/05/2024			086° 00,010' E	SO305/25_3	CTD 25-3			50	
01/05/2024	11:30:00 AM	13° 00,072' N	086° 00,010' E	SO305/25_3	CTD 25-3	SO305_1_1280		300	
01/05/2024	11:30:00 AM	13° 00,072' N	086° 00,010' E	SO305/25_1	CTD 25-1	SO305_1_1270		600	
02/05/2024	3:00:00 PM	12° 59,585' N	087° 59,851' E	SO305/27_4	CTD 27-4	SO305_1_1422		50	
02/05/2024	3:00:00 PM	12° 59,585' N	087° 59,851' E	SO305/27_4	CTD 27-4	SO305_1_1414		300	
02/05/2024	3:00:00 PM	12° 59,585' N	087° 59,851' E	SO305/27_2	CTD 27-2	SO305_1_1402		600	
03/05/2024	2:44:10 AM			SO305/28_2	CTD 28-2	SO305_1_1485		50	
03/05/2024	2:44:10 AM	12° 59,992' N	089° 00,094' E	SO305/28_2	CTD 28-2	SO305_1_1472		300	
03/05/2024	2:44:10 AM	12° 59,992' N	089° 00,094' E	SO305/28_1	CTD 28-1	SO305_1_1465		600	
05/05/2024	11:50:00 PM	15° 00.007 N	85° 45.003 E	SO305/32-2	CTD 32-2	SO305_1_1775		50	Extra
05/05/2024	11:50:00 PM	15° 00.007 N	85° 45.003 E	SO305/32-2	CTD 32-2			300	
05/05/2024	11:50:00 PM	15° 00.007 N	85° 45.003 E	SO305/32-2	CTD 32-2		SO305_1_1759	600	
1/05/2024	3:01:00 AM	16° 00,009' N	089° 36,003' E	SO305/38_6				50	
1/05/2024	3:01:00 AM	16° 00,009' N	089° 36,003' E	SO305/38_6				300	
			089° 36,003' E	SO305/38_6					

5.19 Trace Elements, Trace Element Speciation, Major Ions, Dissolved Organic Carbon and pH

(M. Gledhill¹, A. Firus¹, A. Conventz¹)

¹ GEOMAR

5.19.1 Introduction

Trace elements are essential micronutrients in the ocean, and provide information on material sources and water mass provenance. Many steps in the marine nitrogen cycle, a key focus of SO305, are catalysed by enzymes incorporating trace elements such as iron (Fe), molybdenum (Mo) and copper (Cu). Furthermore, the pronounced oxygen minimum zone in the region can be expected to impact redox sensitive trace elements such as Fe (Moffett et al., 2007; Zhu et al., 2021). Oxygen minimum zones are also characterised by low pH, which influences the chemical equilibria that determines the chemical forms of trace elements present in the water column (chemical speciation) and the binding of trace metals to marine dissolved organic matter (DOM) (Turner et al., 2024).

Rivers and atmospheric deposition are sources of many trace elements to the ocean. The Bay of Bengal receives inputs from the Ganges-Brahmaputra River, which results in a surface low salinity layer and a strongly stratified water column. Furthermore, atmospheric deposition is also significant in the region, with the Bay of Bengal predicted to receive more than 50 μ mol m⁻² yr⁻¹ (Hamilton et al., 2020). The impact of the material sources on the distributions of trace elements in the Bay of Bengal have not been thoroughly studied.

We therefore aimed to characterise the distribution and chemical speciation of the trace elements Fe, Cu, nickel (Ni), zinc (Zn), lead (Pb), cadmium (Cd), manganese (Mn) and cobalt (Co) in the surface waters of the Bay of Bengal. Our aim is to identify the impact of fresh water inputs, atmospheric deposition, physical mixing and internal biogeochemical cycling on trace element distributions in the region.

Despite being among the most abundant elements present in the global ocean, the concentration and distribution of some dissolved major elements (Ca, Sr, Li) are not well studied. Although typically considered conservative, the salinity-normalised concentrations of these major ions do vary by measurable quantities as result of material sources (Steiner et al., 2020). A particular focus of study in this region will be the major ion concentrations in the surface low salinity waters that cap the Bay of Bengal.

5.19.2 Method Description

5.19.2.1 Sample Collection

Near surface water (nominal depth 2.0 ± 0.5 m) was sampled using a trace metal clean towed-fish positioned aft on the starboard side of the ship. Water was pumped directly from the towed-fish through polyethylene tubing into a clean tent supplied with filtered air (MAX 10L HEPA filter) constructed in a laboratory on the SONNE. The towed-fish supply was sampled for dissolved trace metals (DTMs, <0.2 µm), particulate trace metals (PTMs, >0.2 µm), nutrients (nitrate, nitrite, phosphate, silicate), major ions (Ca, Sr, Li and F), pH and dissolved organic carbon (DOC). A

total of 74 towed-fish samples were collected at approximately 3 hourly intervals whilst the ship was underway.

Six depths up to 500 m were sampled from 27 stations using contamination free GoFlo water samplers (General Oceanics) deployed on a Kevlar wire. A mini conductivity-temperature-depth logger (Star-Oddi) was attached to the deepest water sampler. The water samplers were transferred to the clean tent prior to sampling for DTMs, PTMs, dissolved organic trace metals (D-orgTMs), particulate organic trace metals (P-orgTMs), major ions, nutrients (nitrate, nitrite, phosphate, silicate), pH and DOC.

5.19.2.2 Methods

<u>Nutrients</u>: Nutrients were sampled directly (unfiltered) into 15 mL nutrient vials for analysis on board (see section 5.7 above).

<u>pH:</u> pH samples were collected without filtration into 15 mL vials and analysed directly on board via spectrophotometry with the indicator meta-cresol purple using a SAMI-pH logger (Sunburst Sensors). Samples were analysed on the total pH scale at room temperature assuming a salinity of 35. At station 30 the SAMI pH logger developed a technical fault which could not be resolved. pH values will be corrected to in-situ temperature, salinity and pressure post-cruise using the Sunburst Sensor QC-pH routine. pH of surface waters was also logged continuously using a pH logger (Alphox, Pyroscience) deployed in the laboratory at the towed-fish outflow. Data will be available in December 2024.

<u>Major ions:</u> Samples for analysis of major ions were filtered (0.8/0.2 µm, AcroPak 500, Pall) into acid washed 15 mL polypropylene centrifuge tubes and acidified to pH 1.8 with hydrochloric acid. Concentrations of Ca, Sr and Li will be determined at GEOMAR by inductively coupled plasma optical emission spectroscopy (Varian). Data will be available in June 2026.

<u>Dissolved trace metals</u>: Sample for DTMs were filtered (0.8/0.2 µm, AcroPak 500, Pall) into acid washed 125 mL low density polyethylene bottles following GEOTRACES protocols (Cutter et al., 2014). Samples were acidified to pH 1.8 with ultra-high purity hydrochloric acid (Optima, Fisher). Concentrations of iron, manganese, cobalt, copper, cadmium, nickel, lead and zinc will be determined at GEOMAR after preconcentration (SeaFAST, ESI) by high resolution inductively coupled plasma mass spectrometry (Element-XR, Thermo) (Rapp et al., 2017). Data will be available in June 2025.

<u>Dissolved organic carbon</u>: Samples (25 mL) for DOC were filtered (0.8/0.2 µm, AcroPak 500, Pall) into acid washed and pre-combusted (450 °C, 4 hours) 30 mL glass vials sealed with fluorinated ethylene propylene (FEP) septa. Samples were acidified to pH 2 with hydrochloric acid. Dissolved organic carbon and nitrogen will determine at GEOMAR after catalytic oxidation (TOCN-L, Shimadzu). Data will be available in June 2025.

<u>Dissolved and particulate organic trace metals</u>: Samples (ca. 2 L) for D-orgTMs and P-orgTMs were transferred to flexible polyethylene media bags (Flexboy, Sartorius). Water from the bags was passed through a 0.2 µm polyvinylidene fluoride (PVDF) cartridge filter (Sterivax, Millipore) and then over polystyrene divinyl benzene solid phase extraction (SPE) cartridge (500 mg, ENV+, Agilent) using a peristaltic pump at a flow rate of ca. 10 mL/min. Filter and cartridge had been precleaned with methanol (10 mL) and 0.01 M hydrochloric acid (20 mL). Filter cartridges were frozen at -80°C for analysis of the P-orgTM fraction while SPE cartridges were frozen at -20°C

for analysis of D-orgTM fractions. Organic trace metal fractions will be analysed at GEOMAR after extraction from the filter and cartridge by size exclusion and reversed phase high performance liquid chromatography – electrospray ionisation/inductively coupled plasma - mass spectrometry (Thermo) (Gledhill et al., 2022). Data will be available in June 2026.

<u>Particulate trace metals (PTMs)</u>: Particulate trace metals were collected on 0.2 µm acid washed polyethylene sulfone filters (25 mm, PALL) placed in an acid-cleaned polyethylene filter holder and directly attached to the GoFlo bottle tap. GoFlo bottles were pressurized with nitrogen for PTM filtration. For PTM samples collected from the towed-fish, an FEP vacuum unit was connected to the polyethylene filter holder and water was supplied using a flexible media bag (Flexboy, Sartorius). Between 2 and 4 L of seawater was filtered from four depths on each cast. Filters were carefully folded in half, placed into small ziplock bags and frozen at -20°C for transportation back to the laboratory at GEOMAR. Particulate trace metals will be determined after digestion of filters according to GEOTRACES protocols (Al-Hashem et al., 2022; Cutter et al., 2014). Data will be available in June 2026.

Table 5.15 List of samples						
Parameter	Number of GoFlo samples	Number of towed-fish samples				
Nutrients	157	71				
pН	157	73				
Major ions	157	73				
DTMs	157	73				
DOC	157	71				
D-orgTMs	119					
P-orgTMs	119					
PTMs	95	48				

5.19.3 List of Samples

5.19.4 Preliminary Results

Only preliminary pH data are available at the time of writing as most of the analysis of this WP must be undertaken in the laboratory at GEOMAR. A preliminary pH section plot is shown for the latitudinal transect to $14^{\circ}N$ (Fig. 5.12). Oxygen distributions (data: CTD, see section 5.1) are shown for context. pH decreased in the oxygen minimum zone and, in surface waters, towards the north of the transect as the water freshened. The strong relationship between pH and oxygen suggests remineralisation processes made a contribution to decreases in pH in subsurface waters in the Bay of Bengal, although increased CO₂ solubility as a result of lower temperatures can also be expected to be important.

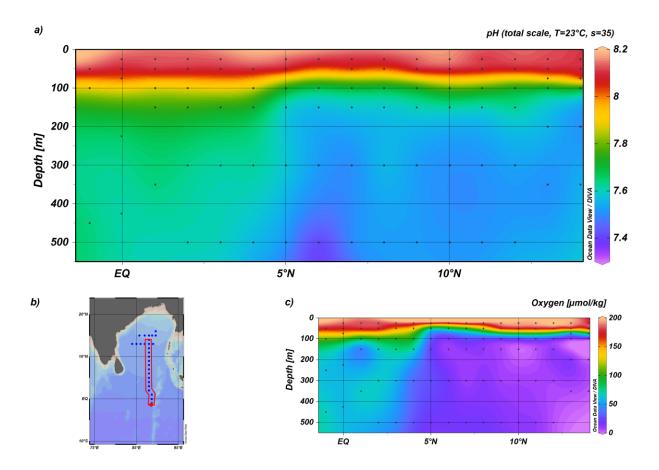


Fig. 5.12 a) Interpolated distribution of pH along a latitudinal transect (1°S, 14°N) at 88°E in the Bay of Bengal.
b) Map showing positions of GoFlo deployment. c) Interpolate oxygen distribution between 1°S and 14 °N at 88°E.

5.20 Nd and REE Isotopes

(A. Conventz¹, M. Gledhill¹, A. Firus¹, E. Hathorne¹, M. Frank¹) ¹ GEOMAR

5.20.1 Introduction

The Bay of Bengal (BoB) is a critical area for studying rare earth elements (REEs) and neodymium (Nd) isotopes due to its unique geochemical and hydrological characteristics. The REE concentrations in the BoB, particularly at various depths, are among the highest in the Indian Ocean (Nozaki et al., 2002). This high concentration is primarily driven by the substantial influx of sediments and freshwater from major rivers, such as the Ganges-Brahmaputra and the Irrawaddy, which collectively contribute a significant amount of terrestrial debris and dissolved materials into the northern part of the Bay (Nozaki et al., 2002).

Nd in these waters reflects the type and age of the rocks in the drainage basins, imparting a distinct geochemical signature to the rivers and seawater through weathering processes (e.g., Goldstein and Jacobsen, 1987; Frank, 2002). The BoB has a thin, low-salinity mixed layer that persists year-round, influenced by substantial freshwater inputs (Sengupta et al., 2006). The massive Bengal and Nicobar fan, the largest submarine fan system on Earth (Curray et al., 2002), also impacts sediment distribution and REE cycling in the region (Hathorne et al., 2020).

Studies have shown elevated REE concentrations in the BoB and Andaman Sea surface waters compared to the rest of the Indian Ocean, with a distinctive REE pattern and unradiogenic Nd isotopes likely resulting from South Asian weathering inputs (Amakawa et al., 2000). These concentrations vary seasonally, correlating with monsoon-induced river discharge (Hathorne et al., 2020).

The enrichment of REEs in the BoB is not solely due to dissolved riverine inputs. The underlying fan sediments act as a sink for dissolved REEs (Nozaki et al., 2002), while the partial dissolution of detrital particles significantly contributes to the dissolved REE pool (Hathorne et al., 2020). These particles are transported by rivers and surface currents, settling through the water column and releasing REEs in the process.

Nd concentration in the BoB surface waters decreases from north to south, influenced by the mixing of radiogenic Indonesian Throughflow waters with less radiogenic BoB waters (Singh et al., 2012). The seasonal monsoon cycle further affects REE distribution, with higher concentrations in surface waters during periods of increased river discharge (Hathorne et al., 2020).

Balancing the Nd budget in the BoB suggests that additional sources, such as release from particulate phases supplied by the Ganges-Brahmaputra river system, are necessary to explain observed distributions. The substantial labile particulate pool, originating from massive river sediment fluxes, plays a crucial role in these seasonal and spatial variations in REE concentrations. Yttrium and REE data also indicate that freshwater input into the BoB can be traced using specific ratios, providing valuable geochemical tracers for monsoon-related freshwater inputs (Yu et al., 2017).

5.20.2 Sample Collection

40 near surface water samples were collected from a towed-fish (nominal depth 2.0 ± 0.5 m) that was positioned on the starboard side of the ship. Through polyethylene tubing the water was pumped into a clean tent constructed in a laboratory on the RV SONNE. In the laboratory, water was filtered (0.8/0.2 µm, AcroPak 500, Pall) into acid cleaned cubitainers (10 L volume) or acid cleaned bottles (125 mL bottles). For later Nd and REE measurements 29 filtered 10 L water samples were collected, including 5 pairs of filtered and unfiltered waters. To increase the resolution additionally 1 L filtered samples with a volume of 125 mL were taken, including one filtered-unfiltered pair, for REE analysis.

From 10 stations 28 samples were collected from GoFlo bottles from at least two depths per station up to 500 m. These 125 mL bottles are again for an increased resolution of REE measurements.

From 3 stations between 8°N and 16°N depth profiles were sampled from a CTD with 10 samples throughout the water column per station. The water was sampled into stable canisters with a volume of 10 L and taken into the clean laboratory. There the samples were filtered with a peristaltic pump over a 142 mm diameter membrane filter within a couple of hours into cubitainers. The membrane filters were folded and saved in plastic bags. Additionally, at a fourth CTD station smaller samples (125 mL) were filtered through AcroPak into small bottles.

All samples were acidified on board with 1 mL concentrated HCl per litre of sample (0.1 %). All further sample preparation for Nd and REE measurements will take place at GEOMAR in Kiel starting in November this year, so measurements will be finished, and data finalized in late summer 2025.

It was planned to take more samples at high resolution in the northern part of the BoB until 18°N, but this was not possible due to an earlier end of the cruise because of a medical emergency.

5.20.3 List of Samples

Table 5.14Sampling at stations (except underway towed-fish sampling between 8°N and 16°N)

Station	Longitude	Latitude	Devices
14	087° 59.962' E	07° 59.971' N	CTD/Ro
15	088° 00.042' E	09° 00.076' N	GoFlo
18	088° 00.283' E	12° 00.239' N	GoFlo
22	085° 04.503' E	12° 44.537' N	GoFlo
23	083° 59.845' E	12° 59.035' N	GoFlo
25	085° 59.889' E	13° 00.025' N	GoFlo
26	086° 59.735' E	12° 59.772' N	GoFlo
27	087° 59.852' E	12° 59.579' N	GoFlo
28	089° 00.094' E	12° 59.988' N	GoFlo
30	087° 59.121' E	13° 59.654' N	GoFlo
32	085° 43.913' E	14° 58.152' N	GoFlo
34	086° 59.932' E	15° 00.115' N	GoFlo
35	087° 59.846' E	15° 00.018' N	GoFlo
36	088° 43.412' E	14° 59.091' N	CTD/Ro
37	089° 38.855' E	15°04.352' N	CTD/Ro
38	089° 35.998' E	15° 59.999' N	CTD/Ro

5.21 Aerosols and Atmospheric Trace Gases

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¹ TROPOS

5.21.1 Introduction

Aerosol particles are important climate drivers as they scatter and absorb solar radiation (direct aerosol effect) and impact the formation and characteristics of clouds (indirect aerosol effect) (Kanakidou et al., 2005). To better understand the diverse climatic effects of aerosol particles, a proper understanding of their sources and transformation/processing is required. However, especially in non-remote marine areas, aerosol particles over the ocean constitute a complex mixture of local marine sources and of non-marine sources with anthropogenic origin (ship emissions) and long-range transported aerosol form continents, which are again a complex mixture of biogenic and anthropogenic sources (Hawkins et al., 2010). In this context, the northern part of the Indian Ocean and in particular the Bay of Bengal (BoB) is the only place in the world where an intense source of continental anthropogenic trace species (and their reaction products) originating from the Northern Hemisphere are directly connected to the pristine air of the Southern Hemisphere by a cross-equatorial monsoonal flow into the Intertropical Convergence Zone (Crutzen and Ramanathan, 2001). Moreover, continental influence is not only driven by

anthropogenic emissions but it can also contain an important fraction of dust. Due to the large content of Iron and Phosphorus on the dust and anthropogenic aerosols, dust deposition can act as fertilizer for the marine biological activities. The strength of continental and anthropogenic emissions on the aerosol particles over the ocean in general, and over northern part of the Indian Ocean in particular, is, however, little understood and poorly-quantified, causing strong uncertainties in the estimation of the climate effect of marine aerosols (Liss and Johnson, 2014).

Despite several research activities carried out in the BoB, detailed knowledge of the atmospheric aerosol chemical composition, sources and processing, especially on the organic fraction and trace metal content of aerosol particles over this region are still very limited. Chemical investigations of aerosol particles were mainly based on offline measurements, which provide no information on the variability of aerosol chemical properties regarding their temporal and spatial resolution as well as information on their mixing state. Altogether, the results from previous campaigns showed that future studies involving a more comprehensive chemical characterization of different aerosol particle sources are essential to better understand the interaction between marine aerosols and long-range transported anthropogenic aerosols and therefore allowing for a proper assessments of Asia's pollution impact on regional and global scales (Balasubramanian et al., 2013; Guazzotti et al., 2003). A scientific understanding of how mixing state affects climate relevant aerosol properties, such as cloud condensation and ice nucleating particle concentrations and aerosol optical properties is still missing over BoB. Consequently, to better understand the sources and the impact to the ocean of the marine boundary aerosol over the BoB, a three-part work approach is deployed in order to better understand: 1) the emissions of primary and secondary aerosol from the marine surface microlayer (SML), 2) the bio-relevance of trace metal on the atmospheric aerosol and their impact on the ocean, and 3) the contribution and mixing state of the different aerosol sources (primary, secondary biogenic or anthropogenic) to the total composition and their impact on climate relevant aerosol properties.

5.21.2 Work at Sea

The atmospheric aerosol measurements were carried out from 16th April 2024 to 11th May 2024 during the expedition. This includes continuous aerosol sampling and measurement activities as well as additional sampling of the SML (sampled from the Zodiac) and sampling from the continuous underway seawater supply. For continuous measurements of physical and chemical aerosol parameters, a laboratory container (Aerosol-Container) equipped with instrumentation was placed on the ninth deck of the ship. Continuous Aerosol-Container measurements include particle number size distributions from 2 nm to 10 µm using a combination of NAIS, SMPS (Scanning Mobility Particle Sizer) and APS (Aerodynamic Particle Sizer), Optical properties, such as absorption and scattering coefficients, were continuously measured by MAAP (Multiangle Absorption Photometer), Aethalometer (AE33), and Nephelometer (Table 5.15). The near realtime measurements of the mixing state and size distribution of airborne BC (black carbon)containing particles were carried out using the Single Particle Soot Photometer (SP2). Further to understand the CCN (cloud condensation nuclei) characteristics, a CCN counter was operated during the cruise. CCN was operational only up to 29th of April. In addition, a standalone CPC was measuring the total aerosol number concentration in 2s time resolution. Together with this, a high-volume filter sampler (PM10) and a Berner impactor (size segregated sampling) was installed

on the roof of the container to collect aerosol particles at a sampling regime of 24 hours during the cruise to provide information on the total particle mass concentration and the chemical composition of the aerosol particles (organic and elemental carbon, OC/EC, water-soluble ions, trace metals etc.). A total of ~ 26 days of filter sampling were carried out during the expedition. During this period, 36 PM10 samples and 24 size segregated sampling was carried out. Together with this several field blanks were also collected to understand the background handling contaminations. A halfback sampler was used to collect the aerosols in the nucleopore filters for the INP (ice nucleating particle) analysis to provide general coverage of INP concentrations throughout the whole cruise. 43 samples were collected during the cruise for the offline INP analysis. Sea water samples were taken regularly throughout the cruise from the onboard underwater pipeline for the analysis of black carbon and INP in the ocean water. All filters and water samples were stored frozen at -20 degrees on the ship for offline analysis at TROPOS after the expedition. Further, an O₃, NO_X, SO₂ analysers were also operated during the cruise to have the concentrations of these gases. Table 5.15 gives an overview of the instrumentation/measurements during the SO305 BIOCAT-IIOE2 campaign.

Instrumentation	Purpose/derived parameters
Digitel DHS-80 with PM10 inlet	Sampling aerosol particles smaller 10 µm
Five stage Berner impactor	Size segregated sampling of aerosol particles with upper cut-off
	diameters of 0.05, 0.14, 0.42, 1.2, 3.5 and 10 μm
Particle sampler (Halfbac)	Sampling aerosol particles for the measurements of INP
	concentration
HR-SP-AMS	Non-refractory aerosol chemical composition (organics, nitrate,
	sulphate, ammonium, chloride) and soot
NAIS/MPSS/APS	Particle number size distribution (PNSD) from 2 nm to 10 μ m.
SP2	Refractory black carbon concentration (rBC), rBC mass size
	distribution, coating thickness on rBC cores.
МААР	Light absorption coefficient at wavelength 635 nm.
Aethalometer	Light absorption coefficients at wavelengths 350, 450, 590, 660,
	880, and 960 nm.
Nephelometer	Light scattering coefficients at wavelength 450, 525 and 635 nm.
CCN counter	CCN concentration
Gas sensors	Ozone, NOx, SO ₂

 Table 5.15
 Overview of the aerosol instrumentation and parameters measured during the SO305 campaign.

A highly advanced high-resolution soot particle aerosol mass spectrometer (HR-SP-AMS) that is a combination of a high-resolution AMS and a single-particle soot photometer (SP2) laser (Onasch et al., 2012; Avery et al., 2020) was deployed during the cruise to study the quantitative chemical information of near PM1 size and mass concentration. This will represent important information to properly apportion the different aerosol particle sources and their physical properties. The HR-SP-AMS is equipped with an intracavity Nd:YAG laser (1064 nm), allowing aerosol particle vaporization of black carbon together with metal (Onasch et al., 2012) together with the determination of the aerosol chemical composition, including total organic matter nitrate, sulphate, ammonium and chloride. With such an instrument, it is possible to perform source apportionment analysis not only on the organic aerosol fraction but also on the refractory black carbon (Bibi et al., 2021; Farley et al., 2022), which will strongly help us to better understand the sources of aerosol over the Bay of Bengal. From the organic matter, MSA mass concentration will be extracted from the AMS data following the method developed by Huang et al. (2018). Source apportionment Advance receptor models will be performed on the dataset by Positive Matrix Factorization (PMF) later. The PMF is a factor analytical method to decompose a matrix of observation (chemical species identified on the filter samples or mass spectra from the AMS) into a set of factor profiles defined by a specific fingerprint and factor contribution (i.e. source contribution to the observed explanatory variable, like total PM mass for the filter samples or organic mass for the AMS). For this purpose, the advanced multilinear engine (Paatero, 1999) approach associated with the source finder program SoFi (Canonaco et al., 2013) will be used. Since our measurements constitutes online and offline aerosol chemical composition, we will take advantage of both the high time resolution provided by the AMS and the comprehensive chemical composition of the offline on a combined source apportionment analysis (Srivastava et al., 2019). Compared to classical source apportionment analysis based on the AMS data, combining online and offline measurements on the same analysis will allow providing chemical information on the AMS PMF factor, which was missing up to now. Parallel to this chemical source apportionment analysis, the PMF approach will also be applied to the MPSS dataset (Ogulei et al., 2007), providing a size-resolved factor.

The originality of our approach is that we applied a broad combination of online and offline aerosol measurements and detailed chemical characterisations associated to seawater and SML chemical studies. Our study will deliver a unique aerosol data set and finally contribute to an advanced understanding of ocean-atmosphere interactions in general and the role of aerosol particles over the BoB in particular. The expected results will be extremely valuable for better modelling particle transport over BoB and the oceans and then improve our prediction on their climate effects.

5.21.3 **Preliminary Results**

Figure 5.13 shows the aerosol particle size distribution measured with SMPS during SO305 BIOCAT IIOE2. We observed the lowest particle concentration during the initial phase of the cruise. These periods were mostly influenced by the precipitation. The aerosol particle size distribution showed bi-modal size distribution with first mode in the Aitken mode (~50 nm) and the second mode in the accumulation mode region (~ 200 nm) during the cruise. Most of the period during the cruise, the accumulation mode was present with higher number concentrations. The total aerosol number shows some periods with very high concentrations. These episodes indicate extra aerosol particles emitted from the exhaust of the research vessel, blown in the direction of the Inlet. These periods will be carefully checked and marked in the final data.

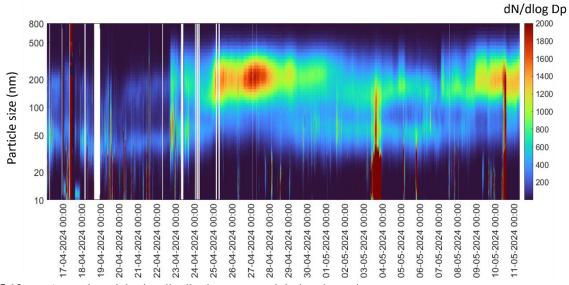


Fig. 5.13 Aerosol particle size distribution measured during the cruise.

Figure 5.14 shows the mass concentrations of different chemical species measured using SP-AMS during the cruise period as well as the daily average concentration of sulphate and ammonium. The sulphate concentration increases as the cruise approaches to the northern most stations. The increased pollutant concentrations mostly refer to transported airmass either from the lands or the sea spray, depending on the wind direction and speed. DMS and sea spray are the main sources of sulphate over the ocean surface.

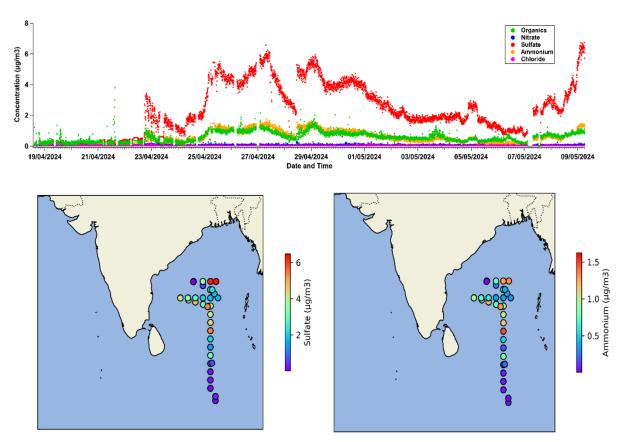


Fig. 5.14 a) Mass concentrations of different chemical species measured during the cruise period, (b-c) shows the daily average concentration of sulphate and ammonium.

6 Station List SO305

6.1 Overall Station List

Station	Device	Date & Time, UTC	Latitude	Longitude	Depth, m
SO305_0_Underway-4	VMADCP_75kHz	2024/04/15 02:30:00	03° 02,115' N	082° 58,218' E	0.0
SO305_0_Underway-5	PS	2024/04/15 02:30:00	03° 02,115' N	082° 58,218' E	0.0
SO305_0_Underway-3	VMADCP_38kHz	2024/04/15 02:30:00	03° 02,115' N	082° 58,218' E	0.0
SO305_0_Underway-6	FBOX	2024/04/15 02:30:00	03° 02,115' N	082° 58,218' E	0.0
SO305_0_Underway-2	EM122	2024/04/15 02:30:00	03° 02,115' N	082° 58,218' E	0.0
SO305_1-1	CTD	2024/04/16 02:59:39	00° 25,947' N	086° 13,861' E	4495.3
SO305_1-2	TMF	2024/04/16 06:28:07	00° 25,946' N	086° 13,890' E	4494.6
SO305_2-1	CTD	2024/04/17 04:22:43	00° 59,971' S	088° 40,211' E	4311.8
SO305_2-2	MSS	2024/04/17 08:26:25	00° 59,992' S	088° 40,211' E	4334.9
SO305_2-3	CTD	2024/04/17 09:25:18	00° 59,932' S	088° 40,217' E	4308.8
SO305_2-5	WS	2024/04/17 11:10:02	00° 59,933' S	088° 40,214' E	4307.8
SO305_2-6	CTD	2024/04/17 13:47:13	00° 59,935' S	088° 40,209' E	4307.2
SO305_3-1	CTD	2024/04/17 18:43:35	00° 30,001' S	088° 40,195' E	4486.8
SO305_4-1	CTD	2024/04/18 00:51:03	00° 01,531' S	088° 40,214' E	4515.8
SO305_4-2	CTD	2024/04/18 04:47:25	00° 01,543' S	088° 40,184' E	4517.7
SO305_4-3	CTD	2024/04/18 05:47:15	00° 01,537' S	088° 40,193' E	4516.5
SO305_4-4	MOOR	2024/04/18 07:22:57	00° 01,458' S	088° 39,926' E	4521.5
SO305_4-5	WS	2024/04/18 15:00:44	00° 00,463' S	088° 39,997' E	4518.4
SO305_4-6	TMF	2024/04/18 16:20:59	00° 00,445' S	088° 40,154' E	4517.9
SO305_5-1	CTD	2024/04/18 19:53:03	00° 29,996' N	088° 40,210' E	4835.7
SO305_5-2	MSS	2024/04/18 23:16:41	00° 30,030' N	088° 40,193' E	4479.4
SO305_6-1	CTD	2024/04/19 04:22:10	01° 00,066' N	088° 40,313' E	4384.7
SO305_6-2	MSS	2024/04/19 07:22:48	01° 00,075' N	088° 40,326' E	4382.0
SO305_6-3	CTD	2024/04/19 08:18:58	00° 59,877' N	088° 39,737' E	4380.6
SO305_6-4	CTD	2024/04/19 09:33:41	00° 59,875' N	088° 39,736' E	4381.2
SO305_6-5	WS	2024/04/19 10:49:07	00° 59,869' N	088° 39,742' E	4380.5
SO305_7-1	CTD	2024/04/19 19:22:07	02° 00,004' N	088° 00,004' E	4281.7
SO305_7-2	MSS	2024/04/19 22:24:02	02° 00,002' N	088° 00,006' E	4279.7
SO305_7-3	WS	2024/04/19 23:17:29	02° 00,025' N	087° 59,891' E	4282.2
SO305_8-1	CTD	2024/04/20 06:45:48	02° 59,998' N	088° 00,002' E	4221.3
SO305_8-2	MSS	2024/04/20 09:46:09	02° 59,999' N	087° 59,998' E	4223.1
SO305_8-3	CTD	2024/04/20 10:34:57	02° 59,782' N	087° 59,736' E	4222.0
SO305_8-4	WS	2024/04/20 11:50:01	02° 59,780' N	087° 59,737' E	4222.8
SO305_9-1	CTD	2024/04/20 19:14:54	03° 59,998' N	088° 00,003' E	4008.8
SO305_9-2	MSS	2024/04/20 22:03:17	03° 59,994' N	087° 59,994' E	4010.7
SO305_9-3	WS	2024/04/20 22:51:14	03° 59,926' N	087° 59,921' E	4007.9
SO305_10-1	CTD	2024/04/21 06:21:51	04° 59,977' N	088° 00,159' E	3976.5
SO305_10-2	DF	2024/04/21 06:43:46	04° 59,974' N	088° 00,157' E	3976.3
SO305_10-3	CTD	2024/04/21 08:45:42	04° 59,996' N	087° 59,998' E	3976.5
SO305_10-4	MSS	2024/04/21 11:32:27	05° 00,000' N	087° 59,999' E	3979.4
SO305_10-5	CTD	2024/04/21 12:35:49	04° 59,996' N	087° 59,914' E	3977.9
SO305_10-6	CTD	2024/04/21 13:54:40	04° 59,987' N	087° 59,915' E	3975.9

SO305_10-7	WS	2024/04/21 16:01:19	04° 59,990' N	087° 59,921' E	3976.5
SO305 10-8	CTD	2024/04/21 18:35:47	04° 59,985' N	087° 59,918' E	3977.2
SO305_10-9	MSS	2024/04/21 19:46:12	04° 59,995' N	087° 59,919' E	3978.3
SO305_10-10	BOAT	2024/04/22 00:26:36	04° 59,997' N	087° 59,911' E	3976.6
SO305_10-11	MSS	2024/04/22 01:37:45	05° 00,271' N	088° 00,130' E	3977.7
SO305_10-12	CTD	2024/04/22 03:01:29	04° 59,927' N	087° 59,779' E	3979.4
SO305_10-13	CTD	2024/04/22 04:31:47	04° 59,926' N	087° 59,774' E	3976.4
SO305_10-14	PUMP	2024/04/22 05:41:25	04° 59,807' N	087° 59,990' E	3976.7
SO305_10-15	MSS	2024/04/22 08:46:06	04° 59,803' N	087° 59,992' E	3977.9
SO305_10-16	CTD	2024/04/22 09:40:38	04° 59,804' N	087° 59,992' E	3978.1
SO305_10-17	FLOAT	2024/04/22 10:49:18	04° 59,804' N	087° 59,995' E	3976.4
SO305_10-18	TMF	2024/04/22 11:01:53	05° 00,184' N	087° 59,835' E	3976.8
SO305_0_Underway-8	uCTD	2024/04/22 12:43:54	05° 14,277' N	087° 59,995' E	3958.9
SO305_11-2	CTD	2024/04/22 18:28:59	05° 59,990' N	088° 00,002' E	3898.3
SO305_11-3	MSS	2024/04/22 21:18:55	05° 59,995' N	087° 59,998' E	3899.3
SO305_11-4	CTD	2024/04/22 22:13:57	05° 59,987' N	088° 00,007' E	3902.4
SO305_11-5	WS	2024/04/22 23:18:42	05° 59,989' N	088° 00,002' E	3898.2
SO305_12-1	DF	2024/04/23 07:11:07	05° 02,717' N	088° 11,104' E	3965.3
SO305_12-2	CTD	2024/04/23 09:30:37	05° 03,350' N	088° 11,803' E	3967.9
SO305_12-3	TMF	2024/04/23 10:22:33	05° 04,287' N	088° 11,712' E	3965.5
SO305_0_Underway-9	uCTD	2024/04/23 11:00:24	05° 07,777' N	088° 11,347' E	3960.0
SO305_13-1	CTD	2024/04/24 01:02:36	07° 00,031' N	087° 59,973' E	3761.9
SO305_13-2	MSS	2024/04/24 01:55:02	07° 00,031' N	087° 59,978' E	3769.2
SO305_13-3	CTD	2024/04/24 02:49:04	07° 00,605' N	088° 00,398' E	3770.1
SO305_13-4	CTD	2024/04/24 06:19:20	07° 00,602' N	088° 00,400' E	3788.7
SO305_13-5	WS	2024/04/24 07:39:34	07° 00,606' N	088° 00,398' E	3775.2
SO305_14-1	CTD	2024/04/24 15:14:50	07° 59,972' N	087° 59,959' E	3658.4
SO305_14-2	MSS	2024/04/24 17:57:09	07° 59,965' N	087° 59,963' E	3657.3
SO305_14-3	CTD	2024/04/24 18:59:05	07° 59,964' N	087° 59,960' E	3656.6
SO305_14-4	WS	2024/04/24 20:18:18	07° 59,971' N	087° 59,958' E	3657.9
SO305_15-1	CTD	2024/04/25 03:36:19	09° 00,075' N	088° 00,043' E	3531.6
SO305_15-2	MSS	2024/04/25 06:23:28	09° 00,072' N	088° 00,048' E	3535.2
SO305_15-3	CTD	2024/04/25 07:27:00	09° 00,077' N	088° 00,043' E	3538.0
SO305_15-4	CTD	2024/04/25 08:49:27	09° 00,075' N	088° 00,047' E	3534.9
SO305_15-5	WS	2024/04/25 10:00:40	09° 00,075' N	088° 00,038' E	3532.3
SO305_16-1	CTD	2024/04/25 17:16:03	10° 00,077' N	088° 00,040' E	3423.3
SO305_16-2	MSS	2024/04/25 19:49:15	10° 00,102' N	088° 00,060' E	3421.5
SO305_16-3	CTD	2024/04/25 20:39:54	10° 00,536' N	088° 00,534' E	3423.5
SO305_16-4	WS	2024/04/25 21:32:37	10° 00,537' N	088° 00,534' E	3422.1
SO305_17-1	MSS	2024/04/26 04:44:37	10° 59,957' N	088° 00,033' E	3332.0
SO305_17-2	CTD	2024/04/26 05:31:02	10° 59,439' N	088° 00,034' E	3335.6
SO305_17-3	WS	2024/04/26 09:27:10	10° 59,477' N	088° 00,046' E	3330.7
SO305_18-1	DF	2024/04/26 16:18:51	11° 58,949' N	087° 59,991' E	3229.7
SO305_18-2	CTD	2024/04/26 18:49:36	11° 59,983' N	087° 59,983' E	3226.7
SO305_18-3	MSS	2024/04/26 21:11:47	11° 59,984' N	087° 59,991' E	3226.4
SO305_18-4	CTD	2024/04/26 21:55:29	12° 00,241' N	088° 00,283' E	3227.5
SO305_18-5	WS	2024/04/26 23:18:52	12° 00,241' N	088° 00,285' E	3224.4

SO305_18-6	BOAT	2024/04/27 00:32:23	12° 00,146' N	087° 59,934' E	3224.3
SO305_18-7	PUMP	2024/04/27 02:50:10	12° 00,144' N	087° 59,939' E	3224.3
SO305_18-7	CTD	2024/04/27 06:32:29	12° 00,143' N	087° 59,937' E	3224.0
SO305_18-9	CTD	2024/04/27 07:40:33	12° 00,139' N	087° 59,933' E	3224.3
SO305_18-10	MSS	2024/04/27 08:51:35	12° 00,139 N 12° 00,188' N	087° 59,977' E	3227.2
SO305_18-11	CTD	2024/04/27 09:43:49	12° 00,501' N	088° 00,220' E	3224.1
SO305_18-11 SO305_18-12	CTD	2024/04/27 11:49:11	12° 00,501' N	088° 00,220 E 088° 00,210' E	3223.9
SO305_18-12 SO305_18-13	CTD	2024/04/27 13:00:47	12° 00,502' N	088° 00,210 E 088° 00,215' E	3223.9
SO305_18-14	MSS	2024/04/27 14:04:10	12° 00,558' N	088° 00,190' E	3223.8
SO305_18-15	CTD	2024/04/27 15:13:37	12° 00,838' N	088° 01,101' E	3192.6
SO305_18-16	CTD	2024/04/27 16:35:49	12° 00,834' N	088° 01,100' E	3225.5
SO305_18-17	MSS	2024/04/27 17:47:36	12° 00,836' N	088° 01,106' E	3223.3
SO305_18-18	FLOAT	2024/04/27 18:40:24	12° 00,919' N	088° 01,502' E	3224.2
SO305_18-19	TMF	2024/04/27 18:42:45	12° 00,933' N	088° 01,531' E	3223.6
SO305_19-1	CTD	2024/04/28 00:55:16	12° 12,330' N	087° 01,567' E	3234.5
SO305_19-2	MSS	2024/04/28 03:28:31	12° 12,322' N	087° 01,556' E	3232.6
SO305_20-1	CTD	2024/04/28 10:28:32	12° 30,009' N	085° 59,988' E	3249.3
SO305 20-2	MSS	2024/04/28 12:49:32	12° 30,008' N	085° 59,990' E	3249.0
SO305_20-3	CTD	2024/04/28 13:40:34	12° 30,153' N	085° 59,579' E	3248.8
SO305_20-4	CTD	2024/04/28 14:54:37	12° 30,154' N	085° 59,583' E	3248.8
SO305_21-1	DF	2024/04/29 01:10:28	11° 56,823' N	087° 37,324' E	3227.7
SO305_21-2	CTD	2024/04/29 03:05:40	11° 56,730' N	087° 37,442' E	3228.2
	TMF	2024/04/29 04:01:36	11° 56,728' N	087° 37,441' E	3227.6
SO305_22-1	CTD	2024/04/29 19:43:07	12° 44,521' N	085° 04,508' E	3255.9
	MSS	2024/04/29 22:11:32	12° 44,535' N	085° 04,503' E	3255.3
SO305_22-3	WS	2024/04/29 23:18:50	12° 44,535' N	085° 04,502' E	3255.9
SO305_23-1	CTD	2024/04/30 07:09:03	13° 00,016' N	084° 00,010' E	3304.8
SO305_23-2	MSS	2024/04/30 09:50:52	12° 59,999' N	084° 00,003' E	3304.1
SO305_23-3	CTD	2024/04/30 10:53:09	12° 59,037' N	083° 59,847' E	3308.3
SO305_23-4	CTD	2024/04/30 12:24:00	12° 59,031' N	083° 59,841' E	3306.5
SO305_23-5	CTD	2024/04/30 13:24:37	12° 59,036' N	083° 59,848' E	3306.8
SO305_23-6	WS	2024/04/30 14:25:52	12° 59,035' N	083° 59,845' E	3307.7
SO305_24-1	CTD	2024/04/30 21:45:42	13° 00,097' N	085° 00,062' E	3238.6
SO305_24-2	MSS	2024/05/01 00:16:53	13° 00,088' N	085° 00,058' E	3238.9
SO305_24-3	CTD	2024/05/01 01:13:24	12° 59,461' N	084° 59,479' E	3238.7
SO305_24-4	CTD	2024/05/01 02:25:07	12° 59,459' N	084° 59,478' E	3240.5
SO305_25-1	CTD	2024/05/01 09:07:44	13° 00,077' N	086° 00,015' E	3199.4
SO305_25-2	MSS	2024/05/01 11:27:23	13° 00,072' N	086° 00,012' E	3200.0
SO305_25-3	CTD	2024/05/01 12:20:51	13° 00,025' N	085° 59,890' E	3199.4
SO305_25-4	CTD	2024/05/01 13:41:53	13° 00,019' N	085° 59,886' E	3198.0
SO305_25-5	WS	2024/05/01 14:30:43	13° 00,025' N	085° 59,889' E	3199.3
SO305_26-1	CTD	2024/05/01 21:38:48	13° 00,097' N	087° 00,002' E	3141.4
SO305_26-2	MSS	2024/05/02 00:05:07	13° 00,094' N	086° 59,996' E	3138.6
SO305_26-3	CTD	2024/05/02 01:04:37	12° 59,776' N	086° 59,729' E	3140.2
SO305_26-4	CTD	2024/05/02 02:08:49	12° 59,775' N	086° 59,738' E	3140.7
SO305_26-5	WS	2024/05/02 03:33:10	12° 59,775' N	086° 59,733' E	3138.3
SO305_27-1	CTD	2024/05/02 10:31:51	13° 00,006' N	088° 00,056' E	3083.6

SO305_27-2	CTD	2024/05/02 10:59:11	12° 59,996' N	088° 00,059' E	3084.5
SO305_27-3	MSS	2024/05/02 13:10:11	12° 59,997' N	088° 00,051' E	3081.0
SO305_27-4	CTD	2024/05/02 14:11:28	12° 59,586' N	087° 59,853' E	3085.7
SO305_27-5	CTD	2024/05/02 15:38:19	12° 59,579' N	087° 59,857' E	3084.2
SO305_27-6	WS	2024/05/02 16:26:01	12° 59,578' N	087° 59,851' E	3083.3
SO305_28-1	CTD	2024/05/02 23:22:09	12° 59,984' N	089° 00,097' E	3041.8
SO305_28-2	CTD	2024/05/03 02:06:34	12° 59,994' N	089° 00,095' E	3040.5
SO305_28-3	WS	2024/05/03 03:23:06	12° 59,989' N	089° 00,094' E	3044.1
SO305_29-1	CTD	2024/05/03 08:44:57	13° 30,657' N	088° 30,724' E	3022.7
SO305_29-2	MSS	2024/05/03 10:59:02	13° 30,657' N	088° 30,711' E	3017.8
SO305_30-1	CTD	2024/05/04 03:01:52	14° 00,000' N	087° 59,998' E	2972.1
SO305_30-2	CTD	2024/05/04 04:21:02	13° 59,998' N	088° 00,001' E	2971.5
SO305_30-3	CTD	2024/05/04 05:42:56	13° 59,999' N	087° 59,999' E	2970.0
SO305_30-4	CTD	2024/05/04 07:35:20	13° 59,998' N	088° 00,003' E	2970.2
SO305_30-5	CTD	2024/05/04 10:00:22	13° 59,996' N	088° 00,004' E	2970.4
SO305_30-6	DF	2024/05/04 12:00:11	14° 00,006' N	088° 00,000' E	2970.3
SO305_30-7	PUMP	2024/05/04 14:14:24	13° 59,650' N	087° 59,123' E	2972.3
SO305_30-8	CTD	2024/05/04 17:21:53	13° 59,655' N	087° 59,124' E	2969.5
SO305_30-9	CTD	2024/05/04 19:13:02	13° 59,643' N	087° 59,125' E	2969.7
SO305_30-10	CTD	2024/05/04 21:59:24	13° 59,646' N	087° 59,124' E	2970.1
SO305_30-11	MSS	2024/05/04 23:11:55	13° 59,649' N	087° 59,123' E	2969.9
SO305_30-12	CTD	2024/05/05 00:03:47	13° 59,653' N	087° 59,119' E	2970.6
SO305_30-13	BOAT	2024/05/05 00:59:47	13° 59,650' N	087° 59,124' E	2972.0
SO305_30-14	WS	2024/05/05 02:37:06	13° 59,643' N	087° 59,122' E	2972.2
SO305_30-15	TMF	2024/05/05 03:31:48	13° 59,643' N	087° 59,122' E	2972.2
SO305_31-1	CTD	2024/05/05 10:21:20	14° 29,934' N	087° 00,037' E	2981.3
SO305_31-2	MSS	2024/05/05 12:42:14	14° 29,942' N	087° 00,023' E	2979.5
SO305_32-1	CTD	2024/05/05 21:31:03	15° 00,025' N	085° 45,032' E	2940.8
SO305_32-2	CTD	2024/05/05 23:11:47	14° 59,996' N	085° 45,007' E	2851.3
SO305_32-3	MSS	2024/05/06 00:01:32	15° 00,001' N	085° 44,999' E	2941.2
SO305_32-4	CTD	2024/05/06 02:24:29	14° 59,439' N	085° 44,731' E	2934.3
SO305_32-5	CTD	2024/05/06 05:31:11	14° 59,434' N	085° 44,738' E	2932.0
SO305_32-6	MSS	2024/05/06 06:33:14	14° 59,405' N	085° 44,726' E	2931.2
<u>SO305_32-7</u>	PUMP	2024/05/06 07:25:24	14° 58,652' N	085° 44,285' E	2936.4
<u>SO305_32-8</u>	CTD	2024/05/06 11:14:13	14° 58,645' N	085° 44,290' E	2938.4
SO305_32-9	MSS	2024/05/06 12:26:27	14° 58,650' N	085° 44,283' E	2937.8
SO305_32-10	CTD	2024/05/06 13:21:49	14° 58,147' N	085° 43,919' E	2937.5
SO305_32-11	CTD WS	2024/05/06 14:44:43	14° 58,151' N	085° 43,920' E	2938.9
SO305_32-12 SO305_32_13	CTD CTD	2024/05/06 15:41:06	14° 58,153' N 14° 58 143' N	085° 43,916' E	2937.0 2935.7
SO305_32-13 SO305_32-14	MSS	2024/05/06 16:53:29 2024/05/06 18:17:24	14° 58,143' N 14° 58,142' N	085° 43,919' E 085° 43,923' E	2935.7 2934.9
SO305_32-14 SO305_32-15	CTD	2024/05/06 19:07:53	14° 57,712' N	085° 43,606' E	2934.9 2934.6
SO305_32-13	DF	2024/05/07 10:48:55	14 57,712 N 13° 59,145' N	083° 43,000 E 088° 12,752' E	2934.6
SO305_33-2	CTD	2024/05/07 12:51:34	13° 59,279' N	088° 12,752 E 088° 13,754' E	2975.0
SO305_33-3	TMF	2024/05/07 13:35:29	13° 59,215' N	088° 13,644' E	2973.0
SO305_33-3	CTD	2024/05/08 05:41:57	15° 00,020' N	086° 59,958' E	2888.5
SO305_34-2	MSS	2024/05/08 08:13:47	15° 00,025' N	086° 59,954' E	2901.0
50505_34-2	100	2027/03/00 00.13.47	15 00,025 N	000 59,954 E	2701.0

SO305_34-3	CTD	2024/05/08 09:07:21	15° 00,120' N	086° 59,932' E	2894.5
SO305_34-4	WS	2024/05/08 10:14:24	15° 00,118' N	086° 59,932' E	2887.4
SO305_35-1	CTD	2024/05/08 16:53:05	15° 00,019' N	087° 59,852' E	2834.3
SO305_35-2	CTD	2024/05/08 18:10:15	15° 00,017' N	087° 59,850' E	2832.9
SO305_35-3	MSS	2024/05/08 20:22:14	15° 00,013' N	087° 59,851' E	2834.9
SO305_35-4	CTD	2024/05/08 21:12:06	15° 00,025' N	087° 59,850' E	2834.4
SO305_35-5	WS	2024/05/08 22:06:25	15° 00,022' N	087° 59,848' E	2832.0
SO305_36-1	CTD	2024/05/09 03:28:32	14° 59,985' N	088° 44,749' E	2844.7
SO305_36-2	MSS	2024/05/09 06:21:39	14° 59,954' N	088° 44,758' E	2843.9
SO305_36-3	CTD	2024/05/09 07:21:44	14° 59,862' N	088° 44,543' E	2847.8
SO305_36-4	CTD	2024/05/09 08:49:26	14° 59,090' N	088° 43,406' E	2841.9
SO305_36-5	WS	2024/05/09 09:58:47	14° 59,093' N	088° 43,405' E	2841.2
SO305_37-1	CTD	2024/05/09 15:56:25	15° 00,016' N	089° 36,018' E	2713.9
SO305_37-2	MSS	2024/05/09 18:03:54	15° 00,016' N	089° 36,022' E	2718.5
SO305_37-3	CTD	2024/05/09 18:55:02	15° 00,145' N	089° 36,139' E	2712.8
SO305_37-4	CTD	2024/05/09 21:27:36	15° 00,142' N	089° 36,135' E	2710.4
SO305_37-5	MSS	2024/05/09 22:46:15	15° 00,142' N	089° 36,133' E	2712.9
SO305_37-6	BOAT	2024/05/09 23:58:09	15° 00,148' N	089° 36,130' E	2713.8
SO305_37-7	PUMP	2024/05/10 02:28:08	15° 00,142' N	089° 36,144' E	2714.6
SO305_37-8	CTD	2024/05/10 06:01:49	15° 02,129' N	089° 37,896' E	2704.8
SO305_37-9	MSS	2024/05/10 07:22:55	15° 02,255' N	089° 37,946' E	2722.7
SO305_37-10	CTD	2024/05/10 08:29:05	15° 04,356' N	089° 38,853' E	2724.6
SO305_37-11	CTD	2024/05/10 10:31:53	15° 04,344' N	089° 38,859' E	2747.4
SO305_37-12	CTD	2024/05/10 11:31:20	15° 04,353' N	089° 38,852' E	2722.4
SO305_37-13	CTD	2024/05/10 13:10:34	15° 04,352' N	089° 38,855' E	2722.8
SO305_37-14	WS	2024/05/10 14:14:18	15° 04,354' N	089° 38,860' E	2723.0
SO305_38-1	WS	2024/05/10 19:57:17	16° 00,018' N	089° 35,980' E	2561.5
SO305_38-2	CTD	2024/05/10 21:02:07	15° 59,998' N	089° 36,004' E	2591.0
SO305_38-3	MSS	2024/05/10 23:10:31	15° 59,999' N	089° 35,998' E	2577.4
SO305_38-4	CTD	2024/05/11 00:08:28	15° 59,998' N	089° 35,999' E	2574.8
SO305_38-5	CTD	2024/05/11 01:12:54	16° 00,001' N	089° 35,993' E	2556.1
SO305_38-6	CTD	2024/05/11 02:20:24	15° 59,998' N	089° 36,000' E	2561.3
SO305_38-7	CTD	2024/05/11 03:20:32	16° 00,008' N	089° 36,006' E	2560.3

Abbreviations:

VMADCP_38kHz: Vessel-mounted ADCP 38 kHz VMADCP_75kHz: Vessel-mounted ADCP 75 kHz PS: Sub bottom profiler FBOX: FerryBox EM122: Multi Beam Echo Sounder EM122 CTD: CTD/Rosette with 22 Bottles TMF: Towed-fish MSS: Microstruture probe WS: GoFlo bottles MOOR: Mooring DF: Drifting particle (sediment) traps BOAT: Zodiac PUMP: Submersible pump FLOAT: Argo float

7 Data and Sample Storage and Availability

The GEOMAR data management group operates the "Ocean Science Information System" (OSIS) as a central information and research data exchange program for marine research projects of GEOMAR and Kiel University. It is publicly accessible and can be used by all cruise participants, including national and international collaborators. OSIS brings together information about expeditions, experiments and numerical models with peer-reviewed publications and available research data. The view of all information in OSIS is publicly accessible, while access to current data in ongoing research projects can be restricted for a definable period of time (moratorium). Alternatively, the submission status of the datasets, including the responsible researcher as contact person, is visible to the public and can promote collaboration with interested researchers. Two GEOMAR data managers are PANGAEA data curators and can advise cruise participants on the preparation of data publication in a "World Data Centre" (e.g. PANGAEA), which then ensures long-term archiving and access to the research data. This data publication process will be based on the files available in OSIS and will therefore be transparent for all reviewers and other researchers. In collaboration with PANGAEA and the "Union for the Application of International Geosample Numbers" (IGSN), the data and samples are made globally traceable, thereby increasing their scientific value and usability. Links to data publishers or PIs provide contact information for external scientists. Seismic, bathymetric and/or hydroacoustic, video and raw image data as well as processed seismic data are archived in GEOMAR's IT storage infrastructure. The processed data will be made publicly available as published data in PANGAEA with the consent of the researcher. Metadata, including contact and access information are provided by OSIS.

The chief scientist and all PIs involved in SO305 BIOCAT-IIOE2 will adhere to the schedule below, which regulates the availability of all information and research data. After SO305 BIOCAT-IIOE2, the GEOMAR data management group will support and guide the researchers in the data sharing and publication process.

- Availability of metadata in OSIS (https://portal.geomar.de/osis): 2 weeks after completion of the cruise and the associated experiments
- Availability of data in OSIS (https://portal.geomar.de/osis):
 12 months after completion of the cruise and associated experiments
- Availability of data in a WDC/PANGAEA (http://www.pangaea.de or as a compilation at http://www.pangaea.de/search?q=campaign:CRUISENAME):
 2 years after completion of the cruise and associated experiments.

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Table 8.1Definition of Data Responsibility for SO305 BIOCAT-IIOE2.

8 Acknowledgements

We acknowledge the financial support of SO305 BIOCAT-IIOE2 by the German Federal Ministry for Education and Research (BMBF) with grants FKZ 03G0305A (GEOMAR) and FKZ 03G0305B (University of Hamburg). TROPOS (grant# HE3086/60-1) and the University of Oldenburg received funding for SO305 from the German Science Foundation (DFG). Hereon (Geesthacht) generously supported the participation of Leon Schmidt in SO305 BIOCAT-IIOE2. Last but not least, the University of Southern Denmark generously supported the participation of the four colleagues from Odense (DK). The Leitstelle Deutsche Forschungsschiffe (German Research Fleet Coordination Centre) at the University of Hamburg provided excellent support during the preparation of SO305 BIOCAT-IIOE2. We thank the Master Tilo Birnbaum and the crew of R/V SONNE for their never-ending excellent support during SO305.

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10 Abbreviations

ADCP stands for Acoustic Doppler Current Profiler.

CTD stands for conductivity, temperature and depth.

Ro stands for rosette ('Kranzwasserschöpfer).

11 Appendices

Table 11.1Underway CTD (uCTD) station list.

Ship station	uCTD station	Date/Time [UTC]	Latitude	Longitude	Speed (kn)
SO305_0_Underway-8	1	22.04.2024, 12:47	05° 14.703' N	087° 59.999' E	8.2
SO305_0_Underway-8	2	22.04.2024, 13:34	05° 21.191' N	088° 00.001' E	8.3
SO305_0_Underway-9	3	23.04.2024, 11:10	05° 09.199' N	088° 11.202' E	8
SO305_0_Underway-9	4	23.04.2024, 13:05	05° 24.402' N	088° 09.665' E	8.1
SO305_0_Underway-9	5	23.04.2024, 14:02	05° 32.087' N	088° 08.888' E	8
SO305_0_Underway-9	6	23.04.2024, 15:07	05° 40.606' N	088° 08.032' E	8
SO305_0_Underway-9	7	23.04.2024, 16:08	05° 48.810' N	088° 07.205' E	7.9
SO305_0_Underway-9	8	23.04.2024, 17:03	05° 56.087' N	088° 06.468' E	7.9
SO305_0_Underway-9	9	23.04.2024, 18:03	06° 04.132' N	088° 05.659' E	8.1
SO305_0_Underway-9	10	23.04.2024, 19:04	06° 12.368' N	088° 04.821' E	8
SO305_0_Underway-9	11	23.04.2024, 20:00	06° 19.817' N	088° 04.059' E	8.2
SO305_0_Underway-9	12	23.04.2024, 21:00	06° 27.997' N	088° 03.238' E	8.1
SO305_0_Underway-9	13	23.04.2024, 22:07	06° 36.971' N	088° 02.328' E	8.2
SO305_0_Underway-9	14	23.04.2024, 23:02	06° 44.533' N	088° 01.561' E	8.1
SO305_0_Underway-9	15	24.04.2024, 00:04	06° 52.766' N	088° 00.728' E	8.1

	Table 11.2	MSS station list.
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Ship Station	MSS Station	Date/Time [UTC]	Latitude	Longitude
SO305 2-2	1	17.04.2024, 08:28	00° 59.993' S	088° 40.211' E
 SO305_5-2	2	18.04.2024, 23:21	00° 30.003' N	088° 40.143' E
SO305 6-2	3	19.04.2024, 07:24	01° 00.073' N	088° 40.321' E
 SO305_7-2	4	19.04.2024, 22:50	02° 00.002' N	087° 59.994' E
 SO305_8-2	5	20.04.2024, 09:48	02° 59.996' N	087° 59.992' E
SO305_9-2	6	20.04.2024, 22:20	03° 59.988' N	087° 59.994' E
SO305_10-4	7	21.04.2024, 11:34	05° 00.001' N	088° 00.005' E
SO305_10-9	8	21.04.2024, 19:49	04° 59.995' N	087° 59.915' E
SO305_10-11	9	22.04.2024, 01:39	05° 00.265' N	088° 00.129' E
SO305_10-15	10	22.04.2024, 08:46	04° 59.802' N	087° 59.992' E
SO305_11-3	11	22.04.2024, 21:19	05° 59.995' N	087° 59.999' E
SO305_13-2	12	24.04.2024, 01:56	07° 00.041' N	087° 59.981' E
SO305_14-2	13	24.04.2024, 17:57	07° 59.965' N	087° 59.963' E
SO305_15-2	14	25.04.2024, 06:23	09° 00.073' N	088° 00.048' E
SO305_16-2	15	25.04.2024, 19:49	10° 00.106' N	088° 00.065' E
SO305_17-1	16	26.04.2024, 04:55	10° 59.966' N	088° 00.044' E
SO305_18-3	17	26.04.2024, 21:12	11° 59.985' N	087° 59.992' E
SO305_18-10	18	27.04.2024, 08:52	12° 00.197' N	087° 59.985' E
SO305_18-14	19	27.04.2024, 14:10	12° 00.573' N	088° 00.231' E
SO305_18-17	20	27.04.2024, 17:51	12° 00.832' N	088° 01.103' E
SO305_19-2	21	28.04.2024, 03:35	12° 12.285' N	087° 01.547' E
SO305_20-2	22	28.04.2024, 12:50	12° 30.010' N	085° 59.985' E
SO305_22-2	23	29.04.2024, 22:13	12° 44.534' N	085° 04.503' E
SO305_23-2	24	30.04.2024, 09:52	12° 59.988' N	084° 00.002' E
SO305_24-2	25	01.05.2024, 00:17	13° 00.080' N	085° 00.051' E
SO305_25-2	26	01.05.2024, 11:28	13° 00.072' N	086° 00.011' E
SO305_26-2	27	02.05.2024, 00:06	13° 00.091' N	086° 59.991' E
SO305_27-3	28	02.05.2024, 13:13	12° 59.992' N	088° 00.046' E
SO305_29-2	29	03.05.2024, 10:59	13° 30.656' N	088° 30.706' E
SO305_30-11	30	04.05.2024, 23:13	13° 59.648' N	087° 59.124' E
SO305_31-2	31	05.05.2024, 12:44	14° 29.935' N	087° 00.011' E
SO305_32-3	32	06.05.2024, 00:02	14° 59.994' N	085° 44.991' E
SO305_32-6	33	06.05.2024, 06:35	14° 59.391' N	085° 44.717' E
SO305_32-9	34	06.05.2024, 12:29	14° 58.645' N	085° 44.280' E
SO305_32-14	35	06.05.2024, 18:17	14° 58.141' N	085° 43.922' E
SO305_34-2	36	08.05.2024, 08:15	15° 00.023' N	086° 59.954' E
SO305_35-3	37	08.05.2024, 20:22	15° 00.013' N	087° 59.852' E
SO305_36-2	38	09.05.2024, 06:24	14° 59.945' N	088° 44.738' E
SO305_37-2	39	09.05.2024, 18:04	15° 00.016' N	089° 36.022' E
SO305_37-5	40	09.05.2024, 22:48	15° 00.143' N	089° 36.132' E
SO305_37-9	41	10.05.2024, 07:49	15° 03.147' N	089° 38.325' E
SO305_38-3	42	10.05.2024, 23:12	15° 59.998' N	089° 35.997' E

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Cast	Ctation.	D-441-	1 - 16				1-4:4-1-	1			1	1	
Cast 2	Station 2.1	Bottle	bedfordn SO305 1	1 0023	depth 4278.2	pressure 4345.8	latitude -1.000	longitude 88.670	year 2024	month 4	day 17	hour 5	minute 48
2	$\frac{2}{2}$ 1	5	SO305 1		2994.1	3032.3	-1.000	88.670	2024	4	17	6	32
2	2 1	7	SO305 1		1500.1	1513.8	-1.000	88.670	2024	4	17	7	7
	$\frac{2}{2}$ 1	8		= /	<u>1500.1</u> 999.2							7	
2	$\frac{2}{2}$ 1	8	SO305 1 SO305 1	1 0030	800.9	1007.1	-1.000	88.670	2024	4	17	7	16
			SO305 1	1 0032 1 0063		806.9		88.670	2024	4	17		21
4	25	2	000000	0000	500.9	504.3	-0.999	88.670	2024		17	14	43
4	25	4	SO305 1	0065	431.4	434.2	-0.999	88.670	2024	4	17	14	51
4	25	11	SO305 1	0072	195.6	196.8	-0.999	88.670	2024	4	17	15	4
4	2 5	13	SO305 1	1 0074	100.5	101.1	-0.999	88.670	2024	4	17	15	14
4	25	14	SO305 1		80.2	80.7	-0.999	88.670	2024	4	17	15	15
4	2 5	15	SO305 1	0076	60	60.3	-0.999	88.670	2024	4	17	15	16
4	25	19	SO305 1	1 0080	50.8	51.1	-0.999	88.670	2024	4	17	15	17
4	2 5	20	SO305 1	0081	30.5	30.7	-0.999	88.670	2024	4	17	15	18
4	2 5	21	SO305 1	1 0082	10.5	10.6	-0.999	88.670	2024	4	17	15	19
6	4 1	4	SO305 1	/	4481.8	4554.7	-0.025	88.670	2024	4	18	2	17
6	4 1	6	SO305 1		3000.1	3038.5	-0.025	88.670	2024	4	18	2	43
6	4 1	7	SO305 1	1 0112	2000.9	2021.7	-0.025	88.670	2024	4	18	3	1
6	4 1	8	SO305 1	1 0113	1500.4	1514.2	-0.025	88.670	2024	4	18	3	9
6	4 1	9	SO305 1	0114	1001.2	1009.2	-0.025	88.670	2024	4	18	3	18
6	4 1	18	SO305 1	1 0123	715.4	720.6	-0.025	88.670	2024	4	18	3	26
6	4 1	22	SO305 1		501.1	504.5	-0.026	88.670	2024	4	18	3	32
7	4 2	7	SO305 1		341.1	343.3	-0.026	88.670	2024	4	18	5	6
7	4 2	10	SO305 1	0137	200.9	202.1	-0.026	88.670	2024	4	18	5	10
7	4 2	14	SO305 1	0141	100.9	101.5	-0.026	88.670	2024	4	18	5	13
7	4 2	16	SO305 1	0143	81.2	81.7	-0.026	88.670	2024	4	18	5	14
7	4 2	21	SO305 1	0143	61.3	61.7	-0.026	88.670	2024	4	18	5	16
7	4 2	22	SO305 1		50.8	51.1	-0.026	88.670	2024	4	18	5	17
8	43	1	SO305 1		31	31.1	-0.026	88.670	2024	4	18	6	33
8	43	6	SO305 1		2.1	2.1	-0.020	88.670	2024	4	18	6	35
10	61	4	SO305 1		4348.8	4418.2	1.001	88.672	2024	4	19	5	47
10	61	5	SO305 1	1 0197	3001.8	3040.1	1.001	88.672	2024	4	19	6	11
10	61	6	SO305 1		2001.8	2021.8	1.001	88.672	2024	4	19	6	28
10	61	0 7	SO305 1 SO305 1		1500.9	1514.7	1.001	88.672	2024	4	19	6	28 38
10	61	8	SO305 1		1000.4	1008.3	1.001	88.672	2024	4	19	6	38 47
10	61	15	SO305 1		759.8	765.4	1.001	88.672	2024	4	19	6	53
10	61	19	SO305 1	0212	501.3	504.7	1.001	88.672	2024	4	19	6	59
10	61	22	SO305 1	0215	370.9	373.3	1.001	88.672	2024	4	19	7	2
11	63	6	SO305 1	0221	200.8	202	0.998	88.662	2024	4	19	8	38
11	63	8	SO305 1	0225	171.2	172.2	0.998	88.662	2024	4	19	8	40
11	63	10	SO305 1		101.3	101.9	0.998	88.662	2024	4	19	8	42
11	63	15	SO305 1	0230	69.6	70	0.998	88.662	2024	4	19	8	44
11	63	22	SO305 1	0257	1.6	1.6	0.998	88.662	2024	4	19	8	48
13	71	1	SO305 1	1 0238	4247.6	4314.4	2.000	88.000	2024	4	19	20	42
13	7 1	3	SO305 1	1 0240	3001.9	3040.3	2.000	88.000	2024	4	19	21	7
13	7 1	4	SO305 1	0241	2000.1	2020.9	2.000	88.000	2024	4	19	21	26
13	7 1	5	SO305 1		1001.4	1009.3	2.000	88.000	2024	4	19	21	45
13	71	9	SO305 1		690.9	695.9	2.000	88.000	2024	4	19	21	54
13	71	11	SO305 1	1 0248	501.7	505.1	2.000	88.000	2024	4	19	21	59
13	7 1	13	SO305 1	1 0250	360.5	362.9	2.000	88.000	2024	4	19	22	3
13	7 1	15	SO305 1	1 0252	200.5	201.7	2.000	88.000	2024	4	19	22	7
13	71	17	SO305 1		101	101.6	2.000	88.000	2024	4	19	22	11
13	71	18	SO305 1		81	81.5	2.000	88.000	2024	4	19	22	12
13	7 1	19	SO305 1		70.3	70.7	2.000	88.000	2024	4	19	22	13
13	71	20	SO305 1		51.4	51.7	2.000	88.000	2024	4	19	22	15
13	7 1	21	SO305 1	0258	31.2	31.3	2.000	88.000	2024	4	19	22	16
13	7 1	22	SO305 1		4.7	4.8	2.000	88.000	2024	4	19	22	18
14	8 1	4	SO305 1	0263	4191.3	4256.7	3.000	88.000	2024	4	20	8	9
14	8 1	5	SO305 1		3000	3038.3	3.000	88.000	2024	4	20	8	30
14	8 1	6	SO305 1		2001	2021.8	3.000	88.000	2024	4	20	8	48
14	8 1	7	SO305 1		1501.2	1515	3.000	88.000	2024	4	20	8	57
14	8 1	8	SO305 1		1000.4	1008.4	3.000	88.000	2024	4	20	9	6
14	8 1	14	SO305 1		759	764.6	3.000	88.000	2024	4	20	9	11
14	8 1	14	SO305 1		498.8	502.1	3.000	88.000	2024	4	20	9	17
14	83	3	SO305 1		339.4	341.6	2.996	87.996	2024	4	20	11	4
15	83	3 7	SO305 1		229.1	230.5	2.996	87.996	2024	4	20	11	8
15	83	9	SO305 1		150.3	151.2	2.996	87.996	2024	4	20	11	8 11
15	83	10	SO305 1 SO305 1	0290	<u>150.3</u> 99.7	100.3	2.996	87.996	2024 2024	4	20	11	11
							2.996	87.996	2024	4		11	
15	83	16	SO305 1		55.1	55.4					20		15
15	83	22	SO305 1		2.4	2.4	2.996	87.996	2024	4	20	11	18
16	91		SO305 1		3978.7	4038.9	4.000	88.000	2024	4	20	20	32
16	91	2	SO305 1		3000.2	3038.6	4.000	88.000	2024	4	20	20	50
16	91	3	SO305 1		2001.8	2022.7	4.000	88.000	2024	4	20	21	8
16	91	4	SO305 1	0307	1501.5	1515.3	4.000	88.000	2024	4	20	21	18
	91	5	SO305 1		1001.1	1009.1	4.000	88.000	2024	4	20	21	28
16	91	9	SO305 1		670.9	675.7	4.000	88.000	2024	4	20	21	36
16	· · ·		00005	0314	500.8	504.2	4.000	88.000	2024	4	20	21	40
16 16	91	11	SO305 1										
16	91 91	15	SO305 1	1 0318	200.7	201.9	4.000	88.000	2024	4	20	21	48
16 16	9 1 9 1 9 1	15 16	SO305 1 SO305 1	1 0318 1 0319	200.7 150.2	201.9 151.1	4.000 4.000	88.000 88.000	2024	4	20	21 21	49
16 16 16	91 91	15	SO305 1	1 0318	200.7	201.9	4.000	88.000				21	

Table 11.3List of samples of halogenated methanes.

16 0 1 0 2 0 0 2 0 2 1 0 1 1 0 1 0 1 0 1 0 1 0 1 0 1 1 0 1 1 0 1 1 1 0 1			1.0	~~~~										
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18 10 3 8 00.3 1 00.3 10.3	18	10 3	4	SO305 1	0329	3943	4002.3	5.000	88.000	2024	4	21	10	3
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36	16 1	19	SO305 1	0740	75	75.4	10.001	88.001	2024	4	25	19	34
36	16 1	21	SO305 1	0742	31.4	31.5	10.001	88.001	2024	4	25	19	37
36	16 1	22	SO305 1	0743	3.7	3.7	10.001	88.001	2024	4	25	19	39
38	17 1	1	SO305	0766	3299.6	3344.7	10.991	88.001	2024	4	26	7	13
38	17 1	2	SO305 1	0767	2000.2	2021.4	10.991	88.001	2024	4	26	7	39
38	17 1	4	SO305 1	0769	1500.3	1514.4	10.991	88.001	2024	4	26	7	50
38	17 1	5	SO305 1	0770	1000	1008.2	10.991	88.001	2024	4	26	8	0
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38	17 1	10	SO305 1	0775	501	504.5	10.991	88.001	2024	4	26	8	
38	1/1	11	SO305 1	0776	400.3	403	10.991	88.001	2024	4	26	8	14
38	17 1	15	SO305 1	0780	200.8	202.1	10.991	88.001	2024	4	26	8	22
38	17 1	18	SO305 1	0783	81.6	82.1	10.991	88.001	2024	4	26	8	27
38	17 1	19	SO305 1	0784	71.7	72.1	10.991	88.001	2024	4	26	8	27
38	17 1	21	SO305 1	0786	31.5	31.7	10.991	88.001	2024	4	26	8	30
38	17 1	22	SO305 1	0787	3.1	3.1	10.991	88.001	2024	4	26	8	32
39	18 2	6	SO305 1	0793	2000.9	2022.1	12.000	88.000	2024	4	26	20	16
39	18 2	10	SO305 1	0797	1493.9	1507.9	12.000	88.000	2024	4	26	20	27
39	18 2	11	SO305 1	0798	1000.9	1009.1	12.000	88.000	2024	4	26	20	36
39	18 2	22	SO305 1	0809	500.7	504.2	12.000	88.000	2024	4	26	20	52
40	18 4	5	SO305 1	0814	442.4	445.4	12.004	88.005	2024	4	26	22	30
40	18 4	7	SO305 1	0816	356.5	358.9	12.004	88.005	2024	4	26	22	33
40	18 4	10	SO305 1	0810	201.2	202.4	12.004	88.005	2024	4	26	22	39
40	18 4	10	SO305	0819	81	202.4 81.5	12.004	88.005	2024	4	26	22	46
40	18 4	15	SO305 1	0824	71.4	71.8	12.004	88.005	2024	4	26	22	47
40	18 4	16	SO305 1	0825	51	51.3	12.001	88.005	2024	4	26	22	49
40	18 4	17	SO305 1	0826	31.1	31.3	12.004	88.005	2024	4	26	22	51
40	18 4	18	SO305 1	0827	11.2	11.2	12.004	88.005	2024	4	26	22	52
40	18 4	22	SO305 1	0831	3.9	3.9	12.004	88.005	2024	4	26	22	54
42	18 9	17	SO305 1	0870	71.2	71.7	12.002	87.999	2024	4	27	8	33
42	18 9	18	SO305 1	0871	50.8	51.1	12.002	87.999	2024	4	27	8	34
42	18 9	19	SO305 1	0872	31	31.2	12.002	87.999	2024	4	27	8	35
42	18 9	20	SO305 1	0873	11.4	11.5	12.002	87.999	2024	4	27	8	36
42	18 9	22	SO305 1	0875	2.2	2.2	12.002	87.999	2024	4	27	8	37
47	18 16	17	SO305 1	0980	73.3	73.7	12.014	88.018	2024	4	27	17	31
47	18 16	18	SO305 1	0981	50.4	50.7	12.014	88.018	2024	4	27	17	32
47	18 16	19	SO305 1	0982	30.8	31	12.014	88.018	2024	4	27	17	33
47	18 16	20	SO305 1	0983	10.7	10.8	12.014	88.018	2024	4	27	17	35
47	18 16	20	SO305	0985	1.7	1.8	12.014	88.018	2024	4	27	17	35
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49	20 1	Í	SO305 1	1008	3216.8	3260.4	12.500	86.000	2024	4	28	11	31
49	20 1	6	SO305 1	1013	2000.1	2021.4	12.500	86.000	2024	4	28	11	53
49	20 1	7	SO305 1	1014	1500	1514.1	12.500	86.000	2024	4	28	12	2
49	20 1	8	SO305 1	1015	1000.5	1008.7	12.500	86.000	2024	4	28	12	11
49	20 1	16	SO305	1023	501.2	504.7	12.500	86.000	2024	4	28	12	23
50	20 2	1	SO305 1	1030	201.6	202.9	12.503	85.993	2024	4	28	13	51
50	20 2	6	SO305 1	1035	171.1	172.1	12.503	85.993	2024	4	28	13	53
50	20 2	9	SO305 1	1038	100.5	101.1	12.503	85.993	2024	4	28	13	57
50	20 2	14	SO305 1	1043	80.7	81.1	12.503	85.993	2024	4	28	13	58
50	20 2	15	SO305 1	1044	50.8	51.1	12.503	85,993	2024	4	28	14	1
50	20 2	16	SO305 1	1045	30.7	30.9	12.503	85.993	2024	4	28	14	2
50	20 2	17	SO305 1	1046	11	11	12.503	85.993	2024	4	28	14	3
50	20 2	22	SO305 1		1.5	1.5	12.503	85.993	2024	4	28	14	4
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53	22 1	13	SO305 1	1108	230.1	231.5	12.742	85.075	2024		29	21	48
53	22 1	17	SO305 1		86.1	86.6	12.742	85.075	2024	4	29	21	53
53	22 1	18	SO305 1	1113	80.9	81.4	12.742	85.075	2024	4	29	21	54
53	22 1	19	SO305 1	1114	50.7	51	12.742	85.075	2024	4	29	21	56
53	22 1	20	SO305 1	1115	30.9	31.1	12.742	85.075	2024	4	29	21	57
53	22 1	21	SO305 1	1116	11.2	11.2	12.742	85.075	2024	4	29	21	59
53	22 1	22	SO305 1	1117	4.5	4.5	12.742	85.075	2024	4	29	22	0
54	23 1	5	SO305 1	1122	3273.9	3318.7	13.000	84.000	2024	4	30	8	21
54	23 1	9	SO305 1	1126	2000.1	2021.4	13.000	84.000	2024	4	30	8	46
54	23 1	10	SO305 1		1500.2	1514.4	13.000	84.000	2024	4	30	8	57
54	23 1	14	SO305 1	1131	1000.5	1008.8	13.000	84.000	2024	4	30	9	7
54	23 1	17	SO305 1	1134	699.8	705	13.000	84.000	2024	4	30	9	15
55	23 3	2	SO305 1	1141	500.4	503.9	12.984	83.997	2024	4	30	11	14
55	23 3	11	SO305 1	1150	249.1	250.7	12.984	83.997	2024	4	30	11	22
55	23 3	14	SO305 1	1153	168.3	169.4	12.984	83.997	2024	4	30	11	26
55	23 3	17	SO305 1	1156	90.6	91.1	12.984	83.997	2024	4	30	11	30
55	23 3	18	SO305 1	1150	80.5	81	12.984	83.997	2024	4	30	11	31
55	23 3	19	SO305 1	1157	55.9	56.2	12.984	83.997	2024	4	30	11	33
55	23 3	20	SO305 1	1158	30.7	30.2	12.984	83.997	2024	4	30	11	35
55	23 3	20	SO305 1		10.9	<u> </u>	12.984	83.997	2024	4	30	11	35
				1160									
55	23 3	22	SO305 1	1161	1.3	1.3	12.984	83.997	2024	4	30	11	37
59	24 3	9	SO305 1	1236	171	172.1	12.991	84.991	2024	5	1	1	31
	24 3	11	SO305 1	1238	83.9	84.4	12.991	84.991	2024	5	1	1	35
59		15	SO305 1	1242	84.4	84.9	12.991	84.991	2024	5	1	1	35
59 59	24 3			1042	50.9	51.2	12.991	84.991	2024	5	1	1	37
59 59 59	24 3	16	SO305 1	1243								-	
59 59		16 17	SO305 1 SO305 1	1243	30.9	30.7	12.991	84.991	2024	5	1	1	39
59 59 59	24 3					30.7 11.5	12.991 12.991	84.991 84.991	2024 2024		1	1	39 40
59 59 59 59	24 3 24 3	17	SO305 1	1244	30.6					5	1 1 1	-	
59 59 59 59 59 59	24 3 24 3 24 3	17 18	SO305 1 SO305 1	1244 1245	30.6 11.4	11.5	12.991	84.991	2024	5 5	1 1 1	1	40
59 59 59 59 59 59 59	24 3 24 3 24 3 24 3	17 18 22	SO305 1 SO305 1 SO305 1	1244 1245 1249	30.6 11.4 3.3	11.5 3.3	12.991 12.991	84.991 84.991	2024 2024	5 5 5	1 1 1 1	1	40 41

61 251 20 50305 1260 124 122 1200 85.998 2024 5 1 12 52 62 253 17 50305 1288 100 81.4 1200 85.998 2024 5 1 12 52 62 253 21 50305 1292 28 100 13000 85.998 2024 5 1 12 58 62 253 21 50305 1292 1290 85.998 2024 5 1 12 5 64 25 4 1290 1209 85.996 2024 5 1 23 65 263 16 50305 1353 13.1 1299 86.996 2024 5 2 1 22 66 263 18 50305 1353 13.0 1299 86.996 2024 5 2 1 2 1 23 </th <th></th> <th>1</th> <th></th> <th></th> <th>1</th>											1			1
62 25.3 15 80305 1284 10.4 122.3 1300 85.998 2024 5 1 12 54. 62 25.3 19 80305 1290 50.7 51 13000 85.998 2024 5 1 12 55. 62 25.3 22 80305 1291 1.9 1.9 1.3000 85.998 2024 5 1 1.3 0 63 63.3 11 80305 1.341 20.0 22.3 1.299 86.996 2024 5 2 1 2.1 1.2 64 63.3 11 81323 7.3.2 1.2996 86.996 2024 5 2 1 2.8 65 63.3 12 80305 1.313 3.8 3.8 12.996 86.996 2024 5 2 1 2.8 2.9 1.3 3.3 3.3 1.3 3.3 1.3 3.3		25 1	19	SO305 1		700.9	706.1	13.001	86.000	2024	5	1	10	59
c2 23 17 S0305 1288 80.9 81.4 13000 85.998 2024 5 1 12 54. 62 25.3 20 S0305 1201 120 11.1 11.00 85.998 2024 5 1 12 58. 63 26.3 12 S0305 1201 1200 1209 85.996 2024 5 2 1 124 65 26.3 11 S0305 1351 59.2 205.5 129.99 85.996 2024 5 2 1 24 66 26.3 17 S0305 1351 59.2 205.5 129.99 85.996 2024 5 2 1 21 25 1 25 1 25 1 23 1 23 24 2024 50.2 1 21 24 1 25 1 23 1 25 1 23 1 25 <td>61</td> <td>25 1</td> <td>20</td> <td>SO305 1</td> <td>1269</td> <td>700.7</td> <td>705.9</td> <td>13.001</td> <td>86.000</td> <td>2024</td> <td>5</td> <td>1</td> <td>10</td> <td>59</td>	61	25 1	20	SO305 1	1269	700.7	705.9	13.001	86.000	2024	5	1	10	59
62 25 3 19 50305 1291 28. 31 3000 85.998 2024 5 1 12 58. 62 25.3 21 50305 1291 13. 11. 11.0 13.000 85.998 2024 5 1 12 58. 62 25.3 21 50305 1291 10.0 10.0 85.998 2024 5 2 1 12 63 63 11 50305 11351 502. 50.7 12.996 86.996 2024 5 2 1 24. 65 63 11 50305 11351 50.2 50.2 12.996 86.996 2024 5 2 1 23. 66 26 11 50305 1331 50.2 50.2 10.20 86.996 2024 5 2 11 23. 66 27 2 50305 1383 30.4 1	62	25 3	15	SO305 1	1286	121.4	122.2	13.000	85.998	2024	5	1	12	52
cc cc s1 13.000 85.998 20.24 5 1 12 56 cc 75.3 21 S0005 1291 28.8 30 13.000 85.998 20.24 5 1 12 58 cc 75.3 21 S0005 1291 13.0 13.00 85.996 20.24 5 2 1 12 13.0 cc 20.3 11.0 S0005 13.83 73.2 12.99 86.996 20.24 5 2 1 23.0 cc 31.3 12.996 86.996 20.24 5 2 1 23.0 cc 31.3 12.996 86.996 20.24 5 2 1 23.0 cc 31.3 12.996 86.996 20.24 5 2 1 23.0 cc 32.000 13.01 13.01 13.00 86.001 20.24 5 2 1 23.0	62	25.3	17	SO305_1	1288	80.9	81.4	13.000	85,998	2024	5	1	12	54
0.2 0.3 0.00 8.5 0.2 0.2 5 1 12 58 0.2 0.3 0.2 1.1 1.000 85.98 20.24 5 1 12 58 0.6 0.5 3 2 0.001 1.901 1.90 1.901 </td <td></td> <td>25.3</td> <td>19</td> <td>SO305_1</td> <td></td> <td></td> <td></td> <td>13,000</td> <td></td> <td></td> <td></td> <td>1</td> <td>12</td> <td></td>		25.3	19	SO305_1				13,000				1	12	
62 23 21 SOUDS 1 [23] 11 11 13 1000 85.998 2024 S 1 13 0 65 26.3 4 SOUDS 1341 1000 12998 80.996 3024 5 2 1 12 66 26.3 16 SOUDS 1343 1001 12996 80.996 2024 5 2 1 1 22 65 26.3 16 SOUDS 1333 132 2 996 80.996 2024 5 2 1 2 65 26.3 18 SOUDS 1339 3.8 12.996 80.996 2024 5 2 2 1 2 1 2 1 2 1 2 1 2 1 2 1 2 1 2 1 2 1 2 1 2 1 2 1 2 1 2												1	12	
62 23 22 320415 1 1 1 0 65 26.3 11 82015 1341 220 223.3 12996 85.098 2024 5 2 1 1 1 24 65 26.3 11 82015 1332 132.7 12996 85.096 2024 5 2 1 1 23 65 26.3 16 80005 1351 10.4 10.5 12996 85.096 2024 5 2 1 2 28 65 26.3 12 80015 1315 10.4 10.5 12996 85.095 2024 5 2 1 1 39 66 74 1 10 80015 10.291 80010 2024 5 2 11 39 66 72 1 80015 1390 2031 12010 88001 2024 5 2												1		
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65 26 3 18 80305 1353 13.4 10.5 12.996 86.996 2024 5 2 1 29 66 26 4 12 80305 1351 38 38 12.996 86.996 2024 5 2 1 15 68 27 2 9 80305 1351 300 85.001 2024 5 2 12 18 68 27 2 9 80305 1401 1304 13000 85.001 2024 5 2 12 21 68 27 2 10 80305 1401 1001 705 13000 88.001 2024 5 2 14 43 69 27 4 17 80305 1421 13.3 14.2 13.9 87.998 2024 5 2 14 45 69 27 4 18 80305 1422 13.0 15.993 87.998 2024 5 2 14 45 69 <td>65</td> <td>26 3</td> <td>16</td> <td>SO305 1</td> <td>1353</td> <td>50.2</td> <td>50.5</td> <td>12.996</td> <td>86.996</td> <td>2024</td> <td>5</td> <td>2</td> <td>1</td> <td>27</td>	65	26 3	16	SO305 1	1353	50.2	50.5	12.996	86.996	2024	5	2	1	27
65 26.3 18 NO305 1353 38 38 38 2996 86.096 2024 5 2 1 29 66 26.4 12 SO305 1353 38 38 12.996 86.096 2024 5 2 2 11 59 68 27.2 9 SO305 1393 13004 13000 88.001 2024 5 2 12 12 68 27.2 9 SO305 1394 13000 88.001 2024 5 2 12 12 68 27.2 10 SO305 1413 330.0 333.1 12.993 87.998 2024 5 2 14 43 69 27.4 19 SO305 1423 33.0 12.993 87.998 2024 5 2 14 45 69 27.4 21 SO305 1424 10.4 10.5 12.993 8	65	26 3	17	SO305 1	1354	31.1	31.3	12.996	86.996	2024	5	2	1	28
65 26 22 80305 137 2024 2037 12996 86095 2024 5 2 2 3 68 27 2 4 80305 1383 3050.5 3090.6 13000 88.001 2024 5 2 2 11 55 68 27 2 3 80305 1393 2000.3 13000 88.001 2024 5 2 12 216 68 27 2 3 80305 1393 1400 88.001 2024 5 2 12 25 68 27 4 10 80305 1421 73.3 72.7 12.993 87.998 2024 5 2 14 44 69 27 4 18 1030 1301 104 10.993 87.998 2024 5 2 14 40 69 27 4 18 100 80.002 </td <td></td> <td>26.3</td> <td>18</td> <td>SO305_1</td> <td>1355</td> <td>10.4</td> <td>10.5</td> <td>12,996</td> <td>86,996</td> <td>2024</td> <td>5</td> <td>2</td> <td>1</td> <td></td>		26.3	18	SO305_1	1355	10.4	10.5	12,996	86,996	2024	5	2	1	
66 26 12 80305 137 12996 86.996 2024 5 2 2 12 18 68 27 2 9 S0305 1385 3050. 13000 88.001 2024 5 2 12 18 68 27 2 13 S0305 1394 150.00 88.001 2024 5 2 12 18 68 27 2 14 S0305 1394 100.00 88.001 2024 5 2 12 44 69 27 4 17 S0305 1421 10.4 10.93 87.998 2024 5 2 14 45 69 27 4 18 S0305 1424 10.4 10.5 12.993 87.998 2024 5 2 14 48 69 27 4 19 S0305 1425 10.0 13.000 80.002 <td< td=""><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td></td><td>1</td><td></td></td<>													1	
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98 35 4 12 SO305 1 2053 83.1 83.6 15.000 87.997 2024 5 8 2 98 35 4 16 SO305 1 2057 53.7 54 15.000 87.997 2024 5 8 2 98 35 4 16 SO305 1 2057 53.7 54 15.000 87.997 2024 5 8 2 98 35 4 17 SO305 1 2058 31.5 31.7 15.000 87.998 2024 5 8 2 98 35 4 18 SO305 1 2059 10.6 10.6 15.000 87.998 2024 5 8 2 98 35 4 19 SO305 1 2060 4.1 4.1 1 5.000 87.998 2024 5 8 2 98 35 4 19 SO305	
98 35 4 16 SO305 1 2057 53.7 54 15.000 87.997 2024 5 8 2 98 35 4 17 SO305 1 2058 31.5 31.7 15.000 87.997 2024 5 8 2 98 35 4 17 SO305 1 2059 10.6 10.6 15.000 87.998 2024 5 8 2 98 35 4 18 SO305 1 2059 10.6 10.6 15.000 87.998 2024 5 8 2 98 35 4 19 SO305 1 2060 4.1 4.1 15.000 87.998 2024 5 8 2	
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102 57 1 17 BOS05 1 2110 50017 50111 11011 05.000 2021 5	19 9
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111 38 4 13 SO305 1 2340 36 36.2 NaN 89.600 2024 5 11 0	0 34
	0 39

Date/Time UTC	Date/Time Local	Number
16.04.2024 12:00	16.04.2024 18:00	30520004
16.04.2024 18:00 17.04.2024 00:00	17.04.2024 00:00	30520010
17.04.2024.06:00	17.04.2024 12:00	30520016
7.04.2024 12:00	17.04.2024 12:00	30520028
7.04.2024 18:00	18.04.2024 00:00	30520034
8.04.2024 00:00	18.04.2024 06:00	30520040
8.04.2024 06:00	18.04.2024 12:00	_30520046
8 04 2024 12:00	18 04 2024 18:00	30520052
8.04.2024 18:00	19.04.2024 00:00	30520058
9 04 2024 00:00	<u>19 04 2024 06:00</u> 10 04 2024 12 00	30520064
<u>9 04 2024 06:00</u> 9.04.2024 12:00	<u> </u>	30520070 30520076
9.04.2024 12:00	20.04.2024 00:00	30520078
0.04.2024.00:00	20.04.2024.06:00	30520088
0.04.2024 06:00	20.04.2024 12:00	30520094
0.04.2024 12:00	20.04.2024 18:00	30520100
0.04.2024.18:00	21.04.2024.00.00	30520106
1.04.2024 00:00	21.04.2024 06:00	_30520112
1.04.2024 06:00	21.04.2024 12:00	30520118
1 04 2024 12:00	21 04 2024 18:00	30520124
2.04.2024.00:00	22.04.2024.06:00	30520136
2.04.2024 06:00	22.04.2024 12:00	30520142 30520148
2.04.2024 12:00	23.04.2024 00:00	30520148
3.04.2024 00:00	23.04.2024 00:00	30520134
3.04.2024 06:00	23.04.2024 12:00	30520166
3 04 2024 12:00	23.04.2024 18:00	30520172
4.04.2024 00:00	24.04.2024 06:00	30520184
4 04 2024 06:00	24.04.2024.12:00	30520190
4.04.2024 12:00	24.04.2024 18:00	30520196
25.04.2024 00:00	25.04.2024 06:00	30520208
25.04.2024 06:00	25.04.2024 12:00	30520214
25 04 2024 12:00 26.04.2024 00:00	<u>25 04 2024 18:00</u> 26.04.2024 06:00	30520220 30520232
6 04 2024 06:00	26.04.2024.06:00	30520232
6 04 2024 12:00	26.04.2024.18:00	30520244
26.04.2024 18:00	27.04.2024 00:00	30520250
27.04.2024 00:00	27.04.2024 06:00	30520256
27 04 2024 06:00	27 04 2024 12:00	30520262
27.04.2024 12:00	27.04.2024 18:00	30520268
28.04.2024 00:00	28.04.2024 06:00	
28 04 2024 06:00	28 04 2024 12:00	30520286
28.04.2024 12:00	28.04.2024 18:00	30520292
29.04.2024 00:00 29.04.2024 06:00	29.04.2024 06:00	30520304 30520310
29 04 2024 12:00	29 04 2024 12:00	30520316
29.04.2024 18:00	30.04.2024 00:00	30520322
30 04 2024 00:00	30.04.2024.06:00	30520328
80.04.2024 03:00	30.04.2024 09:00	30520331
30.04.2024 06:00	30.04.2024 12:00	30520334
80.04.2024 12:00	30.04.2024 18:00	30520340
30 04 2024 18·00	01.05.2024.00:00	30520346
01.05.2024 00:00	01.05.2024 06:00	30520352
01 05 2024 06:00	01 05 2024 12:00	30520358
01.05.2024 12:00	01.05.2024 18:00 02.05.2024 06:00	30520364 30520376
02.05.2024 00:00	02.05.2024 06:00	
2.05.2024 00:00	02.05.2024 12:00	30520388
2.05.2024 18:00	03.05.2024 00:00	30520394
3 05 2024 00:00	03 05 2024 06:00	30520400
3 05 2024 06:00	03 05 2024 12:00	30520406
03.05.2024 12:00	03.05.2024 18:00	30520412
4.05.2024 06:00	04.05.2024 12:00	30520430
4 05 2024 12:00	04 05 2024 18:00	30520436
4.05.2024 18:00	05.05.2024 00:00	30520442
05.05.2024 00:00	05.05.2024 06:00	30520448
05 05 2024 06:00 05.05.2024 12:00	05 05 2024 12:00 05.05.2024 18:00	30520454 30520460
05.05.2024 12:00	05.05.2024 18:00	30520460
06 05 2024 00:00	06.05.2024.06:00	30520400
06.05.2024 06:00	06.05.2024 12:00	30520478
06.05.2024 12:00	06.05.2024 18:00	30520484
07 05 2024 00:00	07.05.2024.06:00	30520496
07.05.2024 06:00	07.05.2024 12:00	30520502
07.05.2024 12:00	07.05.2024 18:00	30520508
08 05 2024 00:00	08 05 2024 06:00	30520520
08.05.2024 06:00	08.05.2024 12:00	30520526
08.05.2024 12:00	08.05.2024 18:00	30520532

 Table 11.4
 List of underway samples of halogenated methanes.

09.05.2024 00:00	09.05.2024 06:00	30520544
09 05 2024 06:00	09.05.2024.12:00	30520550
09.05.2024 12:00	09.05.2024 18:00	30520556
10.05.2024 06:00	10.05.2024 12:00	30520574
10.05.2024 12:00	10.05.2024 18:00	30520580
11 05 2024 07:00	11.05 2024 13:00	30520599

Table 11.5List of underway air samples.

			Canister numb	ers	
		N. 1		D 11	T7: 1
Date/Time UTC	Date/Time Local	Number	014	Double	Kiel
<u>16.04.2024 18:00</u> 17.04.2024 18:00	<u>17.04.2024 00:00</u> 18.04.2024 00:00	<u>30520010</u> 30520034	<u>914</u> 720	956	781 916
18.04.2024 12:00	18.04.2024 00:00	30520054	907	851	677
18.04.2024 12:00	19.04.2024 18:00	30520052	919	631	0//
19.04.2024 13:00	19.04.2024 00.00	30520038	726	802	689/921
19.04.2024 13:00	20.04.2024 00:00	30520082	692	802	009/921
20.04.2024 06:00	20.04.2024 12:00	30520094	548		
20.04.2024 18:00	21.04.2024 00:00	30520106	737	918	945
21.04.2024 09:00	21.04.2024 15:00	30520121	543	710	242
21.04.2024 18:00	22.04.2024 00:00	30520130	992	780	675
22.04.2024 09:00	22.04.2024 15:00	30520145	790		790
22.04.2024 18:00	23.04.2024 00:00	30520154	540		
23.04.2024 13:00	23.04.2024 19:00	30520173	903		
24.04.2024 07:00	24.04.2024 13:00	30520191	763		
24.04.2024 12:00	24.04.2024 18:00	30520196	836		
24.04.2024 18:00	25.04.2024 00:00	30520202	779		
25.04.2024 09:00	25.04.2024 15:00	30520217	535	593	969
25.04.2024 12:00	25.04.2024 18:00	30520220	530		
25.04.2024 18:00	26.04.2024 00:00	30520226	566		
26.04.2024 12:00	26.04.2024 18:00	30520244	799	855	927
26.04.2024 18:00	27.04.2024 00:00	30520250	613		
27.04.2024 07:00	27.04.2024 13:00	30520263	952	958	661
27.04.2024 12:00	27.04.2024 18:00	30520268	718		
27.04.2024 18:00	28.04.2024 00:00	30520274	917		
28.04.2024 06:00	28.04.2024 12:00	30520286	589		
<u>28.04.2024 12:00</u> 28.04.2024 18:00	<u>28.04.2024 18:00</u> 29.04.2024 00:00	<u>30520292</u> 30520298	<u>935</u> 808		
29.04.2024 07:00	29.04.2024 00:00	30520298	744		
29.04.2024 07.00	29.04.2024 13:00	30520311	750		
29.04.2024 12:00	30.04.2024 00:00	30520322	913		
30.04.2024 08:00	30.04.2024 14:00	30520322	567	560	
30.04.2024 13:00	30.04.2024 19:00	30520341	762	500	
30.04.2024 18:00	01.05.2024 00:00	30520346	923		
01.05.2024 09:00	01.05.2024 15:00	30520361	820		
01.05.2024 12:00	01.05.2024 18:00	30520364	685		
01.05.2024 18:00	02.05.2024 00:00	30520370	577	759	544
02.05.2024 08:00	02.05.2024 14:00	30520384	747		
02.05.2024 12:00	02.05.2024 18:00	30520388	782		
02.05.2024 18:00	03.05.2024 00:00	30520394	550		
03.05.2024 09:00	03.05.2024 15:00	30520409	809		
03.05.2024 12:00	03.05.2024 18:00	30520412	978	412	
03.05.2024 23:00	04.05.2024 05:00	30520423	963		
04.05.2024 13:00	04.05.2024 19:00	30520437	940	804	
05.05.2024 07:00	05.05.2024 13:00	30520455	971	791	999
05.05.2024 12:00	05.05.2024 18:00	30520460	766	_	
05.05.2024 18:00	06.05.2024 00:00	30520466	756		
06.05.2024 06:00	06.05.2024 12:00	<u>30520478</u> 30520484	667	+	+
06.05.2024 12:00 06.05.2024 20:00	06.05.2024 18:00 07.05.2024 02:00	30520484 30520492	605 925		1
07.05.2024 07:00	07.05.2024 02:00	30520492	925	-	581
07.05.2024 07.00	07.05.2024 15:00	30520508	857	1	501
08.05.2024 09:00	08.05.2024 15:00	30520508	678	1	1
08.05.2024 09.00	08.05.2024 15:00	30520522	704		
08.05.2024 19:00	09.05.2024 01:00	30520532	646		662
09.05.2024 09:00	09.05.2024 15:00	30520553	973		854
09.05.2024 19:00	10.05.2024 01:00	30520563	694		
10.05.2024 10:00	10.05.2024 16:00	30520578	611		
10.05.2024 17:00	10.05.2024 23:00	30520585	725		942
10.05.2024 18:00	11.05.2024 00:00	30520586	722		<u> </u>
10.05.2024 20:00	11.05.2024 02:00	30520588	946	906	850
11.05.2024 02:00	11.05.2024 08:00	30520594	691	753	949
11.05.2024 06:00	11.05.2024 12:00	30520598	770	795	964
11.05.2024 08:00	11.05.2024 14:00	30520600	658	777	1

Samule Reflord ID Date Time Filtered Samule Bottle Remarks SO285 So285 2004 1.6.41.20 12.0.1 1 More then Infittedion SO285 So285 2005 17.0.1.20 0.000 14.6 5 300 mhar SO385 SO285 10.072 17.0.4.20 21. 1 1 1 SO385 SO385 10.055 1.04.20 1.8.1 3 1 1 SO385 SO385 1.0976 1.7.04.202 1.8.1 3 1 <th>Table 11.</th> <th>6 List of u</th> <th>nderway pig</th> <th>gment sar</th> <th>nples.</th> <th></th> <th></th> <th></th>	Table 11.	6 List of u	nderway pig	gment sar	nples.			
SOURS SOURS 2 0010 16.04.202 18:00 21. 3 second SOURS SOURS 2 0012 17.04.202 06:00 21. 3 Flow-crio A+B zu voll SOURS SOURS 1 0075 17.04.202 18.1 2 Flow-crio A+B zu voll SOURS SOURS 1 0076 17.04.202 1.61. 4 SOURS SOURS 1 0080 17.04.202 1.61. 4 SOURS 2 0040 18.04.202 02.0 1. 6 SOURS 2 0045 18.04.202 1.52.1 SOURS S 0085 1.017 1.84.202 1.1 2.1 SOURS S 0085 1.0151 1.84.202 2.1 4 SOURS S 0085 1.0151 1.84.202 2.1 2 <			Date			Sample Bottle		
SO368 SO368 <th< td=""><td></td><td></td><td></td><td></td><td></td><td>-</td><td>More then 1h filtration</td><td></td></th<>						-	More then 1h filtration	
			16.04.202				200	
SO365 SO365 <th< td=""><td></td><td></td><td></td><td></td><td></td><td></td><td>300 mbar</td><td></td></th<>							300 mbar	
SO206 SO208 SO208 <th< td=""><td></td><td>SO305 1 0072</td><td></td><td>00.00</td><td></td><td></td><td>Flow-cyto A+B zu voll</td><td></td></th<>		SO305 1 0072		00.00			Flow-cyto A+B zu voll	
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SOUBS SOUDS SOUDS <th< td=""><td></td><td></td><td>17.04.202</td><td></td><td></td><td></td><td></td><td></td></th<>			17.04.202					
			17.04.202					
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INDERS INTERNS 7 0136 77 04 707 00:02 71 4	SO305 SO305	SO305 1 0412 SO305 2 0136	<u>22.04.202</u> 22.04.202			6 4		
SO305 SO305 2 0136 22.04.202 00:03 2L 4 SO305 SO305 X 0001 22.04.202 2L Braunglasflasc Zodiac				00:03			Zodiac	
SO305 SO305 A 0001 22.04.202 21 Braungnasnase Zoulat SO305 SO305 1 0428 22.04.202 04:30 2L 1				04:30			200000	
SO305 SO305 1 0429 22.04.202 04:30 2L 2	SO305	SO305 1 0429	22.04.202	04:30	2L			
SO305 SO305 1 0431 22.04.202 04:30 2L 3								
SO305 SO305 1 0432 22.04.202 04:30 2L 4	SO305	50305 1 0432	22.04.202	04:30	2L	4	1	

List of underway nigment sample Tabla 11.6

SO305	SO305 1 0433	22.04.202	04:30	2L	5		
SO305	SO305 1 0434	22.04.202	04:30	2L	6	Fiter lag nicht richtig	lrauf, 50mL weniger
SO305	SO305 2 0142	22.04.202	06:30	2L	1	nach starkem Regen	
SO305	SO305 2 0142 SO305 1 0472	22.04.202	00.50	2L 2L	1	hach starken Kegen	
SO305 SO305	SO305 1 0472 SO305 1 0473	22.04.202		2L 2L	2		
		22.04.202					
SO305	SO305 1 0475	22.04.202		2L	3		
SO305	SO305_1_0476	22.04.202		2L	4		
SO305	SO305 1 0477	22.04.202				Nicht geschlossen	
SO305	SO305 1 0478	22.04.202		2L	6		
SO305	SO305 2 0148	22.04.202	12:00	2L	-		
SO305	SO305 2 0154	22.04.202	18:00	2L	6		
SO305	SO305 2 0194 SO305 1 0506	22.04.202	22:00	2L 2L	1		
SO305	<u>SO305_1_0511</u>	22.04.202	22:00	2L	2		
SO305	SO305 1 0516	22.04.202	22:00	2L	3		
SO305	SO305 1 0517	22.04.202	22:00	2L	4		
SO305	SO305_1_0518	22.04.202	22:00	2L	5		
SO305	SO305 1 0523	22.04.202	22:00	2L	6		
SO305	SO305 2 0160	23.04.202	00:00	2L	-		
SO305	SO305 2 0166	23.04.202	06:00	2L	2		
SO305	SO305 2 0100 SO305 2 0172	23.04.202	12:00	2L 2L	-		
SO305	SO305 2 0172 SO305 2 0184		00:00		-		
		24.04.202		2L			
SO305	<u>SO305_2_0190</u>	24.04.202	06:00	2L	2		
SO305	SO305 1 0592	24.04.202		2L	1	Filter lag nicht richtig	drauf
SO305	SO305 1 0594	24.04.202		2L	2		
SO305	SO305 1 0595	24.04.202		2L	3		
SO305	SO305 1 0596	24.04.202		2L	4		
SO305	SO305 1 0601	24.04.202		2L	5		
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SO305	SO305 2 0196	24.04.202	12:00	2L 2L	-	1	
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SO305	SO305 1 0627	24.04.202	17:00	2L	1		
SO305	SO305 1 0629	24.04.202	17:00	2L	2		
SO305	SO305 1 0630	24.04.202	17:00	2L	3		
SO305	SO305 1 0631	24.04.202	17:00	2L	4		
SO305	SO305 1 0632	24.04.202	17:00	2L	5		
SO305	SO305 1 0633	24.04.202	17:00	2.1L	6		
SO305	SO305 2 0208	25.04.202	00:00	2L	-		
SO305	SO305 2 0200 SO305 2 0214	25.04.202	06:00	2L 2L	5		
SO305	SO305 1 0685	25.04.202	08:00	2L	1		
SO305	SO305 1 0687	25.04.202	08:00	2L	2		
SO305	SO305 1 0688	25.04.202	08:00	2L	3		
SO305	SO305_1_0692	25.04.202	08:00	2L	4	Position dichtet nicht 1	ichtig ab, Disc
SO305	SO305 1 0694	25.04.202	08:00	2L	5		
SO305	SO305 1 0699	25.04.202	08:00	2L	6		
SO305	SO305 2 0220	25.04.202	12:00	$\overline{2L}$	-		
SO305	SO305 1 0736	25.04.202	12:00	2L 2L	1		
SO305	SO305 1 0730 SO305 1 0739	25.04.202	19:00	2L 2L	2		
SO305	SO305_1_0740	25.04.202	19:00	2L	3		
SO305	SO305 1 0741	25.04.202	19:00			Nicht geschlossen	
SO305	SO305 1 0742	25.04.202	19:00	2L	5		
SO305	SO305 1 0743	25.04.202	19:00	2L	6		
SO305	SO305 2 0226	25.04.202	18:00	2L	-		
SO305	SO305 2 0232	26.04.202	00:00	2L	-		
SO305	SO305 2 0238	26.04.202	06:00	2L	2		
SO305	SO305 1 0781	26.04.202	00.00	2L 2L	1		
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SO305	SO305 1 0784	26.04.202		<u>2L</u>	3		l
SO305	SO305 1 0785	26.04.202		2L	4	+	l
SO305	SO305 1 0786	26.04.202		2L	5		
SO305	SO305 1 0787	26.04.202		2L	6		
SO305	SO305_2_0244	26.04.202	12:00	2L	-		
SO305	SO305 2 0250	26.04.202	18:00	2L	-		
SO305	SO305 1 0823	26.04.202	23:00	2L	1		
SO305	SO305 1 0824	26.04.202	23:00	2L	2		
SO305	SO305 1 0825	26.04.202	23:00	2L 2L	3	Flow-cyto C in 129C a	her A+ B in 145
SO305	SO305 1 0825 SO305 1 0826	26.04.202	23:00	2L 2L	4		
	SO305 1 0826 SO305 1 0827		23:00	2L 2L	5	1	
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SO305	SO305 1 0831	26.04.202	23:00	2L	6		l
SO305	SO305 X 0002	27.04.202	02:45	2L	3	Zodiac	
SO305	SO305 1 0869	27.04.202	08:30	2L	1	4	
SO305	SO305 1 0870	27.04.202	08:30	2L	2		
SO305	SO305 1 0871	27.04.202	08:30	2L	3		I
SO305	SO305 1 0872	27.04.202	08:30	2L	4		
SO305	SO305 1 0873	27.04.202	08:30	2L	5		1
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SOURS SOURS FORMS 2704200 18:00 21. 6 Filter sur nich mitize SOURS SOURS 10021 28:044-200 00:00 11. 1 1 SOURS SOURS 10021 28:044-200 00:00 11. 3 Filter nicht mittie SOURS SOURS 10061 28:04-200 00:00 21. 5 1 1 SOURS SOURS 10061 28:04-200 00:00 21. 5 1	SO305	SO305_1_0982	27.04.202	18:00	2L	4	
SOUBS SOUBS <th< td=""><td>SO305</td><td></td><td>27.04.202</td><td>18:00</td><td>2L</td><td>5</td><td></td></th<>	SO305		27.04.202	18:00	2L	5	
SOME SOME <th< td=""><td>SO305</td><td>SO305 1 0985</td><td>27.04.202</td><td>18:00</td><td>2L</td><td>6</td><td></td></th<>	SO305	SO305 1 0985	27.04.202	18:00	2L	6	
SO385 SO385 <th< td=""><td>SO305</td><td>SO305 2 0280</td><td>28.04.202</td><td>00:00</td><td>2L</td><td>-</td><td>Filter war nicht mittig</td></th<>	SO305	SO305 2 0280	28.04.202	00:00	2L	-	Filter war nicht mittig
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SQ38 SQ38 <th< td=""><td></td><td>SO305 1 1003</td><td>28.04.202</td><td>03:00</td><td>2L</td><td>2</td><td>Filter nicht mittig</td></th<>		SO305 1 1003	28.04.202	03:00	2L	2	Filter nicht mittig
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	SO305	SO305 2 0310	29.04.202	06:00	2L	5	filter was dry for short
SQ308 SQ308 <th< td=""><td>SO305</td><td>SO305 2 0316</td><td>29.04.202</td><td>12:00</td><td>2L</td><td>-</td><td></td></th<>	SO305	SO305 2 0316	29.04.202	12:00	2L	-	
SO208 SO2085 SO2085 </td <td>SO305</td> <td>SO305 2 0322</td> <td>29.04.202</td> <td>18:00</td> <td>2L</td> <td>3</td> <td></td>	SO305	SO305 2 0322	29.04.202	18:00	2L	3	
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$\begin{array}{c c c c c c c c c c c c c c c c c c c $							Probe wahrscheinlich unbrauch bar, undicht
	SO305	SO305 1 1159	30.04.202		2L		
$\begin{array}{c c c c c c c c c c c c c c c c c c c $	SO305		30.04.202		2L	5	Druck zwischendurch bei 0,5 bar
$\begin{array}{c c c c c c c c c c c c c c c c c c c $	SO305	SO305 1 1161	30.04.202		2L	6	
$\begin{array}{c c c c c c c c c c c c c c c c c c c $	SO305		30.04.202	18:00	2L	5	
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SO305 Zodiac 10.05.202 02:00 2L Braunglasflasc Zodiac 2m SO305 SO305 1 2212 10.05.202 1 1 1 SO305 SO305 1 2213 10.05.202 2 1 1 SO305 SO305 1 2213 10.05.202 2 2 1 SO305 SO305 1 2214 10.05.202 3 1 1 1 SO305 SO305 1 2215 10.05.202 4 1 1 1 1 SO305 SO305 1 2217 10.05.202 4 1	SO305	SO305 1 2195	09.05.202	22:45	2L	6		
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SO305 SO305 1 2213 10.05.202 2 SO305 SO305 1 2214 10.05.202 3 SO305 SO305 1 2215 10.05.202 4 SO305 SO305 1 2215 10.05.202 4 SO305 SO305 1 2216 10.05.202 5 SO305 SO305 1 2217 10.05.202 6 SO305 SO305 1 2278 10.05.202 6 SO305 SO305 1 2279 10.05.202 12:20 2L SO305 SO305 1 2279 10.05.202 12:20 2L 2 SO305 SO305 1 2279 10.05.202 12:20 2L 3 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2280 10.05.202 12:20 2L 4 leaked, sehr lange gelaufen wegen SO305 SO305 1 2281 10.05.202 12:20 2L 5 sehr lange gelaufen wegen SO305 SO305 1 2282 10.05.202 12:20 2L 5 sehr lange gelaufen wegen SO305 SO305 1 2282 10.	SO305	Zodiac	10.05.202	02:00	2L	Braunglasflasc	Zodiac 2m	
SO305 SO305 1 2214 10.05.202 3 SO305 SO305 1 2214 10.05.202 4 SO305 SO305 1 2215 10.05.202 4 SO305 SO305 1 2216 10.05.202 5 SO305 SO305 1 2217 10.05.202 6 SO305 SO305 1 2278 10.05.202 6 SO305 SO305 1 2279 10.05.202 12:20 SO305 SO305 1 2280 10.05.202 12:20 SO305 SO305 1 2280 10.05.202 12:20 SO305 SO305 1 2281 10.05.202 12:20 2L SO305 SO305 1 2281 10.05.202 12:20 2L 5 So305 SO305 1 2281 10.05.202 12:20 2L 5 SO305 SO305 1 2283 10.05.202 12:20 2L 5 SO305 SO305 1 2283 10.05.202 12:20 <	SO305	SO305 1 2212	10.05.202			1		
S0305 S0305 1 2215 10.05.202 4 S0305 S0305 1 2215 10.05.202 5 S0305 S0305 1 2216 10.05.202 6 S0305 S0305 1 2217 10.05.202 6 S0305 S0305 1 2217 10.05.202 6 S0305 S0305 1 2278 10.05.202 1 S0305 S0305 1 2279 10.05.202 12:20 2L S0305 S0305 1 2279 10.05.202 12:20 2L 3 S0305 S0305 1 2280 10.05.202 12:20 2L 3 sehr lange gelaufen wegen Pumpenproblem S0305 S0305 1 2281 10.05.202 12:20 2L 4 leaked, sehr lange gelaufen wegen S0305 S0305 1 2282 10.05.202 12:20 2L 5 sehr lange gelaufen wegen Pumpenproblem S0305 S0305 1 2283 10.05.202 12:20 2L 6 leaked, sehr lange gelaufen wegen S0305 S0305 2 2032 12:20	SO305	SO305 1 2213	10.05.202			2		
SO305 SO305 1 2216 10.05.202 5 SO305 SO305 1 2217 10.05.202 6 SO305 SO305 1 2217 10.05.202 6 SO305 SO305 1 2278 10.05.202 1 SO305 SO305 1 2279 10.05.202 12:20 2L 1 SO305 SO305 1 2279 10.05.202 12:20 2L 2 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2280 10.05.202 12:20 2L 3 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2280 10.05.202 12:20 2L 4 leaked, sehr lange gelaufen wegen SO305 SO305 1 2281 10.05.202 12:20 2L 5 sehr lange gelaufen wegen SO305 SO305 1 2282 10.05.202 12:20 2L 6 leaked, sehr lange gelaufen wegen SO305 SO305 1 2283 10.05.202 12:20 2L 6 leaked, sehr lange gelaufen wegen SO305 SO305 2 0592 11.05.202 01:45 2L	SO305	SO305 1 2214	10.05.202			3		
SO305 SO305 1 2217 10.05.202 6 SO305 SO305 1 2278 10.05.202 12:20 2L 1 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2279 10.05.202 12:20 2L 2 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2280 10.05.202 12:20 2L 3 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2280 10.05.202 12:20 2L 4 leaked, sehr lange gelaufen wegen SO305 SO305 1 2281 10.05.202 12:20 2L 5 sehr lange gelaufen wegen SO305 SO305 1 2282 10.05.202 12:20 2L 5 sehr lange gelaufen wegen SO305 SO305 1 2282 10.05.202 12:20 2L 6 leaked, sehr lange gelaufen wegen SO305 SO305 1 2283 10.05.202 12:20 2L 6 leaked, sehr lange gelaufen wegen SO305 SO305 2 0592 11.05.202 01:45 2L - SO305 SO305	SO305	SO305 1 2215	10.05.202			4		
SO305 SO305 1 2278 10.05.202 12:20 2L 1 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2279 10.05.202 12:20 2L 2 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2280 10.05.202 12:20 2L 3 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2280 10.05.202 12:20 2L 3 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2281 10.05.202 12:20 2L 4 leaked, sehr lange gelaufen wegen SO305 SO305 1 2282 10.05.202 12:20 2L 5 sehr lange gelaufen wegen SO305 SO305 1 2282 10.05.202 12:20 2L 5 sehr lange gelaufen wegen SO305 SO305 1 2283 10.05.202 12:20 2L 6 leaked, sehr lange gelaufen wegen SO305 SO305 2 0592 11.05.202 01:45 2L - SO305 SO305 1 2338 11.05.202 01:45 2L 1 <	SO305	SO305_1_2216	10.05.202			5		
SO305 SO305 1 2279 10.05.202 12:20 2L 2 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2280 10.05.202 12:20 2L 3 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2281 10.05.202 12:20 2L 4 leaked, sehr lange gelaufen wegen SO305 SO305 1 2282 10.05.202 12:20 2L 5 sehr lange gelaufen wegen SO305 SO305 1 2282 10.05.202 12:20 2L 5 sehr lange gelaufen wegen SO305 SO305 1 2282 10.05.202 1 2:20 2L 5 SO305 SO305 1 2283 10.05.202 1 2:20 2L 6 leaked, sehr lange gelaufen wegen SO305 SO305 2 0592 1 1.05.202 00:00 2L - - SO305 S 0305 1 2338 11.05.202 01:45 2L 1	SO305	SO305 1 2217	10.05.202			6		
SO305 SO305 1 2280 10.05.202 12:20 2L 3 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2281 10.05.202 12:20 2L 4 leaked, sehr lange gelaufen wegen SO305 SO305 1 2282 10.05.202 12:20 2L 5 sehr lange gelaufen wegen SO305 SO305 1 2282 10.05.202 12:20 2L 6 leaked, sehr lange gelaufen wegen SO305 SO305 2 0592 11.05.202 00:00 2L - SO305 SO305 1 0.05.202 01:45 2L 1 SO305 SO305 2 0.338 11.05.202 01:45 2L 1 SO305 SO305 2 0.339 11.05.202 01:45 2L 2	SO305	SO305 1 2278	10.05.202	12:20	2L	1	sehr lange gelaufen we	gen Pumpenproblem
SO305 SO305 1 2281 10.05.202 12:20 2L 4 leaked, sehr lange gelaufen wegen SO305 SO305 1 2282 10.05.202 12:20 2L 5 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2283 10.05.202 12:20 2L 6 leaked, sehr lange gelaufen wegen SO305 SO305 2 0592 11.05.202 00:00 2L -	SO305	SO305 1 2279	10.05.202	12:20	2L	2	sehr lange gelaufen we	gen Pumpenproblem
SO305 SO305 1 2282 10.05.202 12:20 2L 5 sehr lange gelaufen wegen Pumpenproblem SO305 SO305 1 2283 10.05.202 12:20 2L 6 leaked, sehr lange gelaufen wegen SO305 SO305 2 0592 11.05.202 00:00 2L - - SO305 SO305 1 2338 11.05.202 01:45 2L 1 - SO305 SO305 1 2339 11.05.202 01:45 2L 2 -	SO305	SO305 1 2280	10.05.202	12:20	2L	3	sehr lange gelaufen we	gen Pumpenproblem
SO305 SO305 1 2283 10.05.202 12:20 2L 6 leaked, sehr lange gelaufen wegen SO305 SO305 2 0592 11.05.202 00:00 2L - - SO305 SO305 1 2338 11.05.202 01:45 2L 1 - SO305 SO305 1 2338 11.05.202 01:45 2L 1 -	SO305	SO305 1 2281	10.05.202	12:20	2L	4	leaked, sehr lange gela	ufen wegen
SO305 SO305 2 0592 11.05.202 00:00 2L - SO305 SO305 1 2338 11.05.202 01:45 2L 1 SO305 SO305 1 2339 11.05.202 01:45 2L 1	SO305	SO305 1 2282	10.05.202	12:20	2L	5	sehr lange gelaufen we	gen Pumpenproblem
SO305 SO305 1 2338 11.05.202 01:45 2L 1 SO305 SO305 1 2339 11.05.202 01:45 2L 2	SO305	SO305_1_2283	10.05.202	12:20	2L	6	leaked, sehr lange gela	ufen wegen
SO305 SO305 1 2339 11.05.202 01:45 2L 2	SO305	SO305 2 0592	11.05.202	00:00	2L	-		
	SO305	SO305 1 2338	11.05.202	01:45	2L	1		
SO305 SO305 1 2340 11.05.202 01:45 2L 3	SO305	SO305 1 2339	11.05.202	01:45	2L	2		
	SO305	SO305 1 2340	11.05.202	01:45	2L	3		
SO305 SO305 1 2346 11.05.202 01:45 2L 6	SO305	SO305 1 2346	11.05.202	01:45	2L	6		

Date	Time [UTC]	Station	CTD/ TRAP	depth [m]	POC/N, TPC,	mass, micro-	gel traps	meta-g (DNA)	meta-t (RNA),	flow cytom
			cast		PAA, PCHO, lipids, POP, BSi, TEP, CSP	plastics			EEA	•
22.04.24	05:10	SO305/10-13	CTD #1 deploy	5	х					х
22.04.24	05:10	SO305/10-13	Cast 023	50	х			х	х	х
22.04.24	05:10	SO305/10-13		100	х					х
22.04.24	05:10	SO305/10-13		150	х					х
22.04.24	05:10	SO305/10-13		200	х					х
22.04.24	05:10	SO305/10-13		300	х			х	х	х
22.04.24	05:10	SO305/10-13		400	х					х
22.04.24	05:10	SO305/10-13		500	х					х
22.04.24	05:10	SO305/10-13		600	х			х	x	х
23.04.24	09:31	SO305/12-2	CTD #1 magazien	5	х					x
23.04.24	09:31	SO305/12-2	#1 recover Cast 027	50	x			x	х	x
23.04.24	09:31	SO305/12-2		100	х					x
23.04.24	09:31	SO305/12-2		150	х					x
23.04.24	09:31	SO305/12-2		200	х			x	х	x
23.04.24	09:31	SO305/12-2		300	х					х
23.04.24	09:31	SO305/12-2		400	x					x
23.04.24	09:31	SO305/12-2		500	x			х	x	x
23.04.24	09:31	SO305/12-2		600	x					x
23.04.24	08:48	SO305/12-1	TRAP #1	50	x	X	x		x	
23.04.24	08:48	SO305/12-1		100	x	x	х			
23.04.24	08:48	SO305/12-1		150	x	x	x			
23.04.24	08:48	SO305/12-1		200	x	x	x			
23.04.24	08:48	SO305/12-1		300	x	x				
23.04.24	08:48	SO305/12-1		400	x	х	х			
23.04.24	08:48	SO305/12-1		500	x	х	х			
23.04.24	08:48	SO305/12-1		600	x	х			x	
27.04.24	07:40	SO305/18-8	CTD	5	x					x
27.04.24	07:40	SO305/18-8	#2 deploy Cast 041	50	x			x	x	x
27.04.24	07:40	SO305/18-8		100	x					x
27.04.24	07:40	SO305/18-8		150	x					x
27.04.24	07:40	SO305/18-8		200	x			x	x	x
27.04.24	07:40	SO305/18-8		300	x					x
27.04.24	07:40	SO305/18-8		400	x					x
27.04.24	07:40	SO305/18-8		500	x			x	x	x
27.04.24	07:40	SO305/18-8		600	x					x
29.04.24	04:00	SO305/21-2	CTD	5	x					x
29.04.24	04:00	SO305/21-2	#2 recover Cast 052	50	x			x	x	x
29.04.24	04:00	SO305/21-2		100	x					x
29.04.24	04:00	SO305/21-2		150	х					х

Table 11.7List of samples from drifting sediment traps.

29.04.24 04:00 SO305/21-2 200 x x x x 29.04.24 04:00 SO305/21-2 300 x x x x 29.04.24 04:00 SO305/21-2 400 x x x x 29.04.24 04:00 SO305/21-2 400 x x x x 29.04.24 04:00 SO305/21-2 500 x x x x 29.04.24 04:00 SO305/21-2 600 x x x x 29.04.24 04:00 SO305/21-2 600 x x x x 29.04.24 08:48 SO305/21-1 TRAP#2 50 x x x x 29.04.24 08:48 SO305/21-1 100 x x x x 29.04.24 08:48 SO305/21-1 150 x x x x 29.04.24 08:48 SO305/21-1 200 x x x x 29.04.24 08:48 SO	x x x x x x
29.04.24 04:00 SO305/21-2 400 x x x x x 29.04.24 04:00 SO305/21-2 500 x x x x x 29.04.24 04:00 SO305/21-2 600 x x x x x 29.04.24 08:48 SO305/21-2 500 x x x x 29.04.24 08:48 SO305/21-1 TRAP#2 50 x x x x 29.04.24 08:48 SO305/21-1 TRAP#2 50 x x x x 29.04.24 08:48 SO305/21-1 100 x x x x 29.04.24 08:48 SO305/21-1 150 x x x x 29.04.24 08:48 SO305/21-1 200 x x x x	x x
29.04.24 04:00 SO305/21-2 500 x x x x 29.04.24 04:00 SO305/21-2 600 x x x x 29.04.24 08:48 SO305/21-1 TRAP #2 50 x x x x 29.04.24 08:48 SO305/21-1 TRAP #2 50 x x x x 29.04.24 08:48 SO305/21-1 100 x x x x 29.04.24 08:48 SO305/21-1 150 x x x x 29.04.24 08:48 SO305/21-1 150 x x x x 29.04.24 08:48 SO305/21-1 200 x x x x	x
29.04.24 04:00 SO305/21-2 600 x 29.04.24 08:48 SO305/21-1 TRAP #2 50 x x x 29.04.24 08:48 SO305/21-1 TRAP #2 50 x x x 29.04.24 08:48 SO305/21-1 100 x x x x 29.04.24 08:48 SO305/21-1 150 x x x x 29.04.24 08:48 SO305/21-1 200 x x x x	
29.04.24 08:48 SO305/21-1 TRAP #2 50 x x x x 29.04.24 08:48 SO305/21-1 100 x x x x 29.04.24 08:48 SO305/21-1 100 x x x x 29.04.24 08:48 SO305/21-1 150 x x x 29.04.24 08:48 SO305/21-1 200 x x x	x
29.04.24 08:48 SO305/21-1 100 x x x x 29.04.24 08:48 SO305/21-1 150 x x x x 29.04.24 08:48 SO305/21-1 150 x x x x 29.04.24 08:48 SO305/21-1 200 x x x x	
29.04.24 08:48 SO305/21-1 150 x x x 29.04.24 08:48 SO305/21-1 200 x x x	
29.04.24 08:48 SO305/21-1 200 x x x	
29.04.24 08:48 SO305/21-1 300 x x x	
29.04.24 08:48 SO305/21-1 400 x x x	
29.04.24 08:48 SO305/21-1 500 x x x	
29.04.24 08:48 SO305/21-1 600 x x x	
05.05.24 00:05 SO305/30-12 CTD 5 x	х
05.05.24 00:05 SO305/30-12 #3 deploy Cast 082 50 x x x x	х
05.05.24 00:05 SO305/30-12 100 x	x
05.05.24 00:05 SO305/30-12 150 x	х
05.05.24 00:05 SO305/30-12 200 x	х
05.05.24 00:05 SO305/30-12 300 x x x	х
05.05.24 00:05 SO305/30-12 400 x	х
05.05.24 00:05 SO305/30-12 500 x	x
05.05.24 00:05 SO305/30-12 600 x x x x	x
05.05.24 23:50 SO305/32-2 Extra CTD 5 x	x
05.05.24 23:50 SO305/32-2 1 24h St. 50 x x x	x
05 05 24 23:50 SO305/32-2 no trap 100 v	x
05.05.24 23:50 SO305/32-2 Cast 085 100 X 05.05.24 23:50 SO305/32-2 150 X	х
05.05.24 23:50 SO305/32-2 200 x x x x	x
05.05.24 23:50 SO305/32-2 300 x	x
05.05.24 23:50 SO305/32-2 400 x	x
05.05.24 23:50 SO305/32-2 500 x x x	x
05.05.24 23:50 SO305/32-2 600 x	x
07.05.24 12:51 SO305/33-2 CTD 5 x	x
#3 recover	x
07.05.24 12:51 SO305/33-2 Cast 093 50 x x x x x 07.05.24 12:51 SO305/33-2 100 x	x
07.05.24 12:51 SO305/35-2 150 X	x
	x x
	X
07.05.24 12:51 SO305/33-2 500 x x x	x
07.05.24 12:51 80305/33.2 600 *	x
07.05.24 12:51 SO305/33-2 600 x	1
07.05.24 11:05 SO305/33-1 TRAP #3 50 x x x x	
07.05.24 11:05 SO305/33-1 TRAP #3 50 x x x x 07.05.24 11:05 SO305/33-1 TRAP #3 100 x x x x	
07.05.24 11:05 SO305/33-1 TRAP #3 50 x x x x	

07.05.24	11:05	SO305/33-1		300	х	х			х	
07.05.24	11:05	SO305/33-1		400	х	х	х			
07.05.24	11:05	SO305/33-1		500	х	х	x			
07.05.24	11:05	SO305/33-1		600	х	х			х	
11.05.24	02:45	SO305/38-6	Extra CTD	5	Х					х
11.05.24	02:45	SO305/38-6	2 no trap	50	х			х	х	х
11.05.24	02:45	SO305/38-6	Cast 113	100	х					х
11.05.24	02:45	SO305/38-6		150	х					х
11.05.24	02:45	SO305/38-6		200	х					х
11.05.24	02:45	SO305/38-6		300	х			х	x	х
11.05.24	02:45	SO305/38-6		400	х					х
11.05.24	02:45	SO305/38-6		500	х					х
11.05.24	02:45	SO305/38-6		600	Х			x	х	x