

Preliminary Report
of
The Hakuho Maru Cruise KH-86-3

June 3 - August 1, 1986
The Bering Sea and
the northern North Pacific

Ocean Research Institute
University of Tokyo
1986

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By
The Scientific Members of the Expeditions
Edited by
Akihiko HATTORI
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Preface

This volume compiles the oceanographic data obtained during the KH-86-3 cruise of the R/V Hakuho Maru from 3 June to 1 August 1986. Included also are summaries of the individual research work carried out by the scientists aboard.

During the last decade, several cooperative oceanographic research efforts have been made in the Bering Sea and the northern North Pacific aimed at the advancement of our understanding of physical, chemical and biological processes, and of their interaction in the most productive sea area in the world. Within the framework of this cooperative research, two Hakuho Maru cruises were conducted in 1975 and 1978. The present cruise formed the third phase of this integrated research program, and had close connections with the ISHTAR program sponsored by the Office of Polar Program, the National Science Foundation, USA. The data obtained in the earlier cruises appear in the reports of the Hakuho Maru Cruises KH-75-4 and KH-78-3, Ocean Research Institute, University of Tokyo, 1977 and 1979.

On behalf of the scientists aboard, I wish to express my sincere gratitude to Captain I. Tadama, the other officers and the crew members of the Hakuho Maru for their cooperation throughout the cruise, and to Mr. Toshisuke Nakai and Ms. Masae Otsu for editorial assistance.

1 December 1986

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Chief Scientist
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Outline of the cruise

The cruise consisted of three legs: leg 1, from Tokyo to Kodiak, Alaska; leg 2, from Kodiak to Vancouver, British Columbia; and from Vancouver to Tokyo (Fig. 1). The location of the oceanographic stations and the dates are given in Table 1.

Emphasis was placed on chemical, biochemical and biological aspects with special reference to cycling of biophilic elements. Detailed information was collected for : (1) distribution of temperature and salinity, (2) distribution of dissolved gases, nutrients, chlorophylls, trace metals, dissolved and particulate organic matter, and stable and radioactive nuclides, (3) distribution of phytoplankton, zooplankton and micronekton, and (4) underwater irradiance and turbidity. Simultaneous shipboard experiments on carbon and nitrogen metabolism were carried out.

At selected stations, bottom sediment samples were collected with a box corer, and settling materials by installing sediment traps at various depths. Gases and aerosols in maritime air were also collected over the extended areas of the northern North Pacific throughout the cruise.

The names and institutions of the scientists who participated in this cruise are listed in Table 2, and observation items at each station in Table 3.

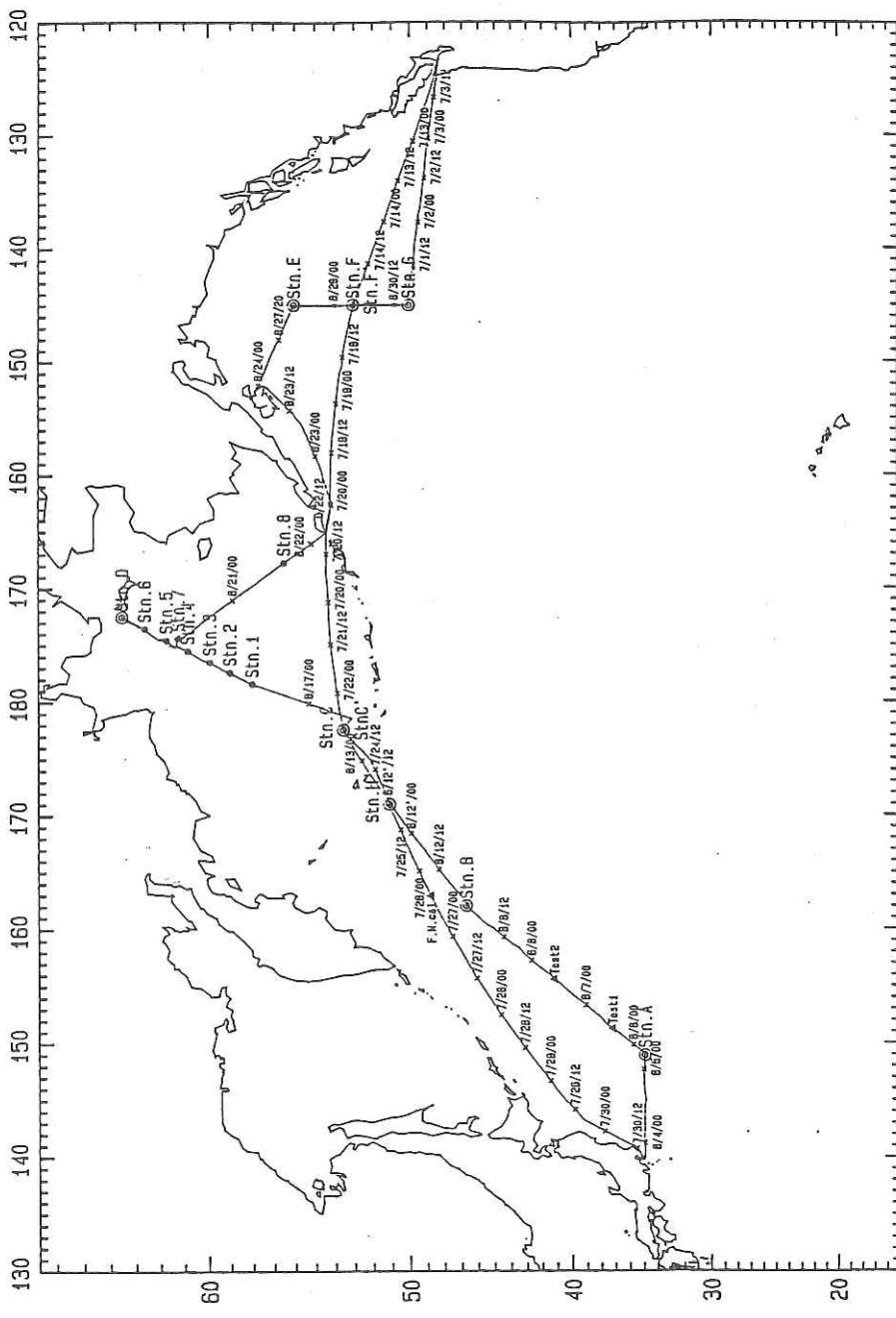


Fig. 1. Track chart of the KH-86-3 cruise of the Hakuho Maru.

Table 1. Location of oceanographic stations and dates

Leg	Station	Latitude	Longitude	Date
Leave Tokyo				6/ 3/86
1 A	34° 48.9'N-34° 57.3'N	148° 50.7'E-148° 57.7'E		6/ 5/86
B	46° 38.0'N-46° 44.2'N	160° 01.5'E-160° 30.4'E		6/ 9/86-6/12/86
C	52° 58.0'N-53° 32.3'N	177° 30.2'E-178° 33.7'E		6/13/86-6/16/86
1	57° 58.9'N-57° 59.9'N	178° 24.8'W-178° 27.8'W		6/17/86
2	58° 59.5'N-59° 00.0'N	177° 24.2'W-177° 28.1'W		6/18/86
3	59° 53.9'N-59° 55.0'N	176° 28.8'W-176° 31.5'W		6/18/86
4	60° 48.7'N-60° 49.3'N	175° 29.6'W-175° 30.2'W		6/18/86
5	61° 42.5'N-61° 42.8'N	174° 32.1'W-174° 32.2'W		6/18/86-6/19/86
6	62° 35.8'N-62° 35.9'N	173° 31.9'W-173° 32.4'W		6/19/86
D	63° 30.0'N-63° 31.1'N	172° 29.3'W-172° 30.0'W		6/19/86
7-1	61° 12.7'N-61° 13.0'N	174° 19.8'W-174° 20.4'W		6/20/86
7-2	61° 10.9'N	174° 19.3'W		6/20/86
7-3	61° 08.6'N	174° 18.6'W		6/20/86
7-4	61° 05.0'N	174° 27.0'W-174° 27.1'W		6/20/86
8	56° 30.2'N-56° 30.7'N	167° 44.9'W-167° 47.2'W		6/21/86
Arrive Kodiak, Alaska				6/24/86
Leave Kodiak, Alaska				6/27/86
2 E	55° 59.9'N-56° 01.3'N	145° 00.4'W-145° 03.6'W		6/28/86
F	52° 59.4'N-53° 03.9'N	144° 57.1'W-145° 00.3'W		6/29/86-6/30/86
G	49° 59.8'N-50° 03.8'N	144° 56.7'W-144° 59.9'W		6/30/86-7/ 1/86
Arrive Vancouver, British Columbia				7/ 4/86
Leave Vancouver, British Columbia				7/11/86
3 F'	52° 55.7'N-53° 01.3'N	144° 38.6'W-144° 59.3'W		7/14/86-7/17/86
C'	53° 29.1'N-53° 30.7'N	177° 28.5'E-177° 32.3'E		7/22/86
H	51° 01.2'N-51° 05.4'N	170° 56.4'E-170° 59.6'E		7/24/86-7/25/86
Arrive Tokyo				8/ 1/86

Table 2. Scientists aboard

Akihiko HATTORI	Ocean Research Insititute, University of Tokyo
Isao KOIKE	Ocean Research Insititute, University of Tokyo
Makoto TERAZAKI	Ocean Research Insititute, University of Tokyo
Toshisuke NAKAI	Ocean Research Insititute, University of Tokyo
Toshiro SAINO	Ocean Research Insititute, University of Tokyo
Ken FURUYA	Ocean Research Insititute, University of Tokyo
Hirotaka OTOBE	Ocean Research Insititute, University of Tokyo
Masae OTSU	Ocean Research Insititute, University of Tokyo
Hae-Lip SUH	Ocean Research Insititute, University of Tokyo
Luis TUPAS	Ocean Research Insititute, University of Tokyo
Shizuo TSUNOGAI	Faculty of Fisheries, Hokkaido University
Koh HARADA	Faculty of Fisheries, Hokkaido University
Kazunori TAGUCHI	Faculty of Fisheries, Hokkaido University
Seiya NAGAO	Faculty of Fisheries, Hokkaido University
Kouji DEMURA	Faculty of Fisheries, Hokkaido University
Kuninao TADA	Faculty of Fisheries, Hokkaido University
Tsuneo ODATE	Faculty of Fisheries, Hokkaido University
Satoru KANAMORI	Water Research Institute, Nagoya University
Masahiko YAMATO	Water Research Institute, Nagoya University
Akio HASHIMOTO	Water Research Institute, Nagoya University
Eiichiro TANOUE	Environmental Science, the Graduate School of Science and Technology, Kobe Univesity
Shigemitsu HARA	Environmental Science, the Graduate School of Science and Technology, Kobe Univesity
Hiroshi SASAKI	Faculty of Agriculture, Tohoku University
Haruto ISHII	Faculty of Agriculture, Tohoku University
Nobuyuki KADOYA	Faculty of Marine Science and Technology, Tokai University
Tatsuo AONO	Faculty of Science and Technology, Kinki University
Masashi KUSAKABE	Department of Geological Science, University of Southern California
Tadashi YOSHINARI	New York State Department of Health

Table 3. Observation Items at each station

Station	A	B	C	1	2	3	4	5	6	D	<u>7</u>	8	E	F	G	F'	C'	H
	1	2	3	4	3	4	5	6	7	8	1	2	3	4				
CTD (+RMSx24) observation	x	x	x								x	x	x	x	x	x	x	x
OCTOPUS (+RMSx12) observation	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x
Water sampling																		
Niskin	x	x	x							x								
Large Volume	x	x									x							x
Clean	x	x	x							x	x					x	x	
Van Dorn	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	
Sediment sampling																		
Box Corer	x	x	x							x						x	x	
OKEAN																x	x	
Plankton sampling																		
NORPAC	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x	x
ORI	x										x	x	x	x	x	x	x	x
MTD	x	x	x	x											x	x	x	
ORI-MPS	x														x	x	x	
Sediment trap															x	x	x	
Anchored	x	x	x													x	x	
Floating	x	x													x	x	x	

Routine observations of oceanographic variables

At each station, casts of CTD fitted with an oxygen electrode and twenty four 5-liter rosette water samplers were made to collect information on water temperature, salinity, dissolved oxygen, phosphate, silicic acid, nitrate, nitrite, ammonium, pH, alkalinity, chlorophylls, and particulate matter. Additional water samples were collected with 23-liter Niskin bottles or 20-liter van Dorn bottles from the upper 200 m layer. Casts of OCTOPUS (Ishimaru et al., 1984) were also conducted in the shallow layer (< 200 m), and data on underwater irradiance, in situ chlorophyll fluorescence, and turbidity were simultaneously collected. Saino was responsible for the operation of CTD and OCTOPUS. The names of the persons who carried out the measurements are given after each item.

Water temperature was measured using a pair of protected reversing thermometers (Nakai), and salinity with a Guildline Autosal 8400A salinometer (Otobe, Kusakabe, Tada and Odate). Dissolved oxygen was determined by the Winkler method adopting a Hirama photometric end-point detector (Tsunogai and Saino). The obtained values for temperature, salinity and oxygen were used for calibration of CTD sensors (Nakai).

Silicic acid, nitrate, nitrite and ammonium were determined using a Technicon autoanalyzer AAII (Hattori, Otsu and Saino). The methods described in Strickland and Parsons (1972) were used with modifications for autoanalyzer measurements. Phosphate was determined manually by the method of Murphy and Riley (Harada, Taguchi and Nagao). The pH was determined with a precision of 0.003 pH unit using a specially designed pH meter at 25 °C, and the alkalinity from the pH shift after the addition of a definite amount of HCl to the seawater samples (Kanamori, Hashimoto, Kadoya and Aono, cf. page 72). Total dissolved nitrogen was measured using a Yanako TN-7 total nitrogen analyzer (Koike and Tupas, cf. page 71).

Particulate matter was collected on a Whatman type F glass fiber filter, and chlorophyll *a* and phaeophytin were determined fluorometrically by the method described in Strickland and Parsons (1972) (Furuya, Tanoue and Hara). Numbers and volumes of particulate matter were determined with an Elzone 80XY particle counter (Sasaki and Ishii).

The data available at present are summarized in Tables 4-37, and Figs. 2-10.

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- Murphy, J. and J.P. Riley (1962). A modified single solution method for the determination of phosphate in natural waters. *Anal. Chim. Acta*, 27, 31-36.
- Strickland, J.D.H. and T.R. Parsons (1972). A Practical Handbook of Seawater Analysis. Fisheries Research Board of Canada, Ottawa.

Table 4. Summary of hydrographic data at Station B

Station	B			Date			June 9, 1966			Lat.			46° - 39.9 N			Air T.			5.9 C			Weather			Fog			Sea					
	Depth	5630 m	TIME	02 : 40 - 05 : 46	Long.	162° - 01.9 E	Barrow.	1005.6 mb	Wind	NE-2.0 m/s	Swell	3				CTD data	D	T	S	D.O.	Bst	Debt	C/L/t	m/l	c	m/l	c	m/l	c	m/l	c		
Sample No.	D	T	S	D.O.	pH	AT	S102	P04	N03	N02	NH4	Pot-T	Sig-t	%DO	AOU	D	T	S	D.O.	Bst	Debt	C/L/t	m/l	c	m/l	c	m/l	c	m/l	c	m/l	c	
AtS or HYDROCAST	m	°C	m/l	mg/l			um	um	um	um	um			%		m/l	m/l	m/l	m/l	m/l	m/l	m/l	m/l	m/l	c	m/l	c	m/l	c	m/l	c	m/l	c
0	0	4.11	33.013	7.65	7.863	2.311	39.8	1.90	22.6	0.21	0.39	4.110	26.19	104.2	-0.31	0	4.107	33.022	7.74	180.5	0.0000												
50	50	2.96	33.053	7.72	7.839	2.309	40.9	1.98	23.7	0.21	0.46	2.957	26.33	102.3	-0.17	10	4.109	33.024	7.77	180.5	0.0189												
100	101	1.24	33.166	7.48	7.776	2.323	45.4	2.02	27.1	0.40	0.72	1.235	26.56	94.9	0.40	20	3.837	33.020	7.77	178.1	0.0364												
200	200	2.96	33.795	1.76	7.410	2.397	92.8	3.10	43.9	0.02	0.98	2.948	26.93	23.4	5.75	30	3.741	33.019	7.79	177.3	0.0540												
300	300	3.33	33.993	0.77	7.366	2.388	106	3.28	45.3	0.00	0.14	3.310	27.05	10.4	6.66	50	3.543	33.030	7.76	174.7	0.0895												
400	400	3.40	34.132	0.49	7.376	2.401	118	3.27	46.3	0.00	0.36	3.373	27.16	6.6	6.92	75	1.729	33.126	7.53	153.2	0.1304												
500	500	3.25	34.218	0.48	7.389	2.439	127	3.28	45.7	0.00	0.25	3.216	27.24	6.5	6.96	100	1.360	33.159	7.29	148.2	0.1690												
600	601	3.10	34.280	0.48	7.399	2.429	131	3.23	46.0	0.00	0.40	3.060	27.30	6.4	6.98	150	2.582	33.622	3.67	121.7	0.2365												
700	701	2.94	34.327	0.50	7.407	2.466	141	3.25	45.7	0.00	0.26	2.893	27.35	6.7	6.99	200	2.971	33.816	2.13	110.2	0.2945												
800	800	2.80	34.361	0.54	7.417	2.453	144	3.21	45.7	0.00	0.14	2.747	27.39	7.2	6.97	300	3.168	34.003	0.99	97.8	0.3998												
900	901	2.67	34.394	0.58	7.424	2.447	152	3.19	44.6	0.00	0.10	2.610	27.43	7.7	6.96	400	3.411	34.139	0.51	69.6	0.4961												
1000	1001	2.54	34.427	0.70	7.435	2.455	155	3.17	45.0	0.00	0.17	2.474	27.47	9.3	6.96	500	3.282	34.224	0.51	81.9	0.5851												
1250	1249	2.26	34.496	0.97	7.462	2.453	162	3.11	44.4	0.00	0.22	2.497	27.55	12.8	6.64	600	3.089	34.290	0.47	75.5	0.6676												
1500	1500	2.08	34.546	1.27	7.495	2.472	166	3.09	43.7	0.00	0.19	1.980	27.60	16.6	6.37	700	2.942	34.339	0.51	70.4	0.7450												
1750	1750	1.92	34.584	1.54	7.528	2.482	167	3.02	43.1	0.00	0.19	1.802	27.64	20.1	6.13	800	2.782	34.377	0.55	66.3	0.8175												
2000	1999	1.80	34.613	1.94	7.555	2.467	167	2.92	42.0	0.00	0.18	1.663	27.68	24.8	5.78	1000	2.487	34.449	0.66	58.4	0.9525												
2250	2250	1.69	34.635	2.30	7.590	2.475	162	2.86	40.3	0.00	0.11	1.533	27.70	29.8	5.41	1250	2.257	34.511	0.92	51.8	1.1046												
2500	2500	1.60	34.663	2.60	7.615	2.477	160	2.77	39.9	0.00	0.06	1.422	27.72	33.6	5.13	1500	2.067	34.559	1.17	46.7	1.2439												
2750	2750	1.55	34.664	2.85	7.638	2.479	158	2.64	39.4	0.00	0.18	1.350	27.74	36.8	4.89	1750	1.907	34.597	1.49	42.6	1.3730												
3000	3000	1.51	34.672	3.06	7.655	2.488	155	2.66	38.1	0.00	0.08	1.286	27.75	39.5	4.69	2000	1.784	34.625	1.88	39.6	1.4942												
3250	3250	1.48	34.677	3.22	7.670	2.485	153	2.61	36.4	0.00	0.06	1.232	27.75	41.5	4.53	2250	1.681	34.646	2.28	37.3	1.6102												
3500	3498	1.46	34.683	3.28	7.677	2.476	152	2.60	38.1	0.00	0.22	1.186	27.76	42.3	4.48	2500	1.602	34.661	2.57	35.5	1.7219												
3750	3751	1.45	34.685	3.49	7.686	2.470	150	2.47	37.1	0.00	0.16	1.149	27.76	45.0	4.27	2750	1.552	34.672	2.83	34.4	1.8309												
4000	4000	1.45	34.689	3.56	7.695	2.464	148	2.54	37.3	0.00	0.38	1.121	27.76	45.9	4.20	3000	1.509	34.680	3.07	33.4	1.9391												
4500	4353	1.49	34.686	3.65	---	---	145	---	36.6	0.00	0.03	1.119	27.76	47.1	4.10	3250	1.479	34.687	3.24	32.7	2.0457												
5000	4843	1.53	34.690	3.70	---	---	145	---	37.2	0.00	0.12	1.097	27.76	47.8	4.04	3500	1.463	34.692	3.40	32.3	2.1527												
5500	5322	1.58	34.690	3.76	---	---	145	---	36.6	0.00	0.05	1.083	27.76	48.6	3.97	4000	1.446	34.698	3.64	31.7	2.3681												

Table 5. Summary of hydrographic data at Station C

Station	C	Date	June 13, 1986			Lat.			53° - 30.5 N			Ain T.			5.5 C			Weather			Cloudy			Sea		
			Depth	3720 m	TIME	13:58	- 17:05	Long.	177	- 38.6 E.	Barrow	1006.3 mb	Wind	W	- 7.5 m/s	Swell	1	Dist	0.0	Dist	0.0	Dist	0.0	Dist	0.0	
No. S or HYDROCAST		D	T	S	D.O.	pH	AT	met/l	PO4	N03	NH4	Pot-T	Sig-t	%DO	AOU	D	T	S	D.O.	D	T	S	D.O.	D	T	S
No.	m	°C						um	um	um	um	um	um	%	ml/l	um	°C	%	ml/l	um	°C	%	ml/l	um	°C	%
0	0	5.14	33.113	7.67	7.892	2.308	31.2	1.79	20.5	0.17	0.50	5.140	26.16	107.2	-0.52	0	5.142	33.122	7.25	183.5	0.0000					
50	50	4.41	33.156	7.60	7.864	2.311	34.1	1.86	21.5	0.20	0.53	4.406	26.28	104.4	-0.32	10	5.044	33.110	7.17	183.3	0.0196					
100	100	3.53	33.268	6.00	7.739	2.305	48.9	2.14	29.4	0.20	0.20	3.523	26.45	80.7	1.43	20	4.888	33.122	7.20	180.4	0.0379					
200	200	3.72	33.509	4.37	7.618	2.316	65.9	2.53	34.6	0.00	0.15	3.706	26.63	59.2	3.02	30	4.567	33.140	7.17	176.1	0.0552					
300	300	3.83	33.796	2.42	7.494	2.336	81.0	2.87	38.9	0.00	0.09	3.809	26.85	32.9	4.93	50	4.334	33.138	7.13	173.9	0.0905					
400	400	3.66	33.936	1.35	7.422	2.354	95.6	3.07	43.9	0.00	0.09	3.692	26.97	18.3	6.03	75	3.713	33.217	6.85	162.0	0.1324					
500	500	3.54	34.041	0.96	7.402	2.382	108	3.15	45.2	0.00	0.05	3.505	27.07	13.0	6.43	100	3.458	33.319	6.05	152.0	0.1720					
600	600	3.47	34.108	0.85	7.401	2.383	114	3.18	43.9	0.00	0.10	3.428	27.13	11.5	6.55	150	3.417	33.427	5.39	143.6	0.2466					
700	701	3.36	34.170	0.84	7.403	2.385	122	3.19	44.5	0.00	0.10	3.311	27.19	11.3	6.58	200	3.770	33.561	4.46	136.6	0.3465					
800	801	3.24	34.230	0.64	7.409	2.402	128	3.13	45.7	0.00	0.08	3.184	27.25	8.6	6.80	300	3.835	33.590	2.56	149.9	0.4471					
900	901	3.11	34.279	0.62	7.412	2.407	138	3.18	45.4	0.00	0.04	3.047	27.30	8.3	6.84	400	3.719	33.619	1.63	109.1	0.5642					
1000	1001	3.00	34.317	0.61	7.416	2.407	140	3.18	45.7	0.00	0.00	2.930	27.34	8.2	6.87	500	3.568	34.027	1.07	99.6	0.6719					
1250	1251	2.70	34.463	0.79	7.429	2.437	152	3.17	45.7	0.00	0.00	2.613	27.46	10.5	6.74	600	3.468	34.114	0.78	92.2	0.7716					
1500	1500	2.39	34.477	0.87	7.449	2.445	168	3.14	44.7	0.00	0.13	2.286	27.52	11.5	6.72	700	3.352	34.188	0.64	85.4	0.8652					
1750	1751	2.12	34.539	1.07	7.475	2.462	177	3.11	43.1	0.00	0.10	1.999	27.59	14.0	6.57	800	3.236	34.244	0.56	80.2	0.9831					
2000	2001	1.92	34.583	1.36	7.563	2.468	194	3.04	43.1	0.00	0.06	1.784	27.64	17.7	6.31	1000	2.996	34.330	0.51	71.5	1.1166					
2250	2251	1.78	34.613	1.62	7.533	2.470	200	3.03	40.4	0.00	0.03	1.621	27.68	21.0	6.08	1250	2.684	34.419	0.55	62.2	1.3008					
2500	2501	1.70	34.634	1.90	7.565	2.473	205	2.97	40.9	0.00	0.01	1.520	27.70	24.6	5.81	1500	2.369	34.493	0.70	54.0	1.4850					
2750	2751	1.64	34.649	2.03	7.573	2.472	210	2.83	40.1	0.00	0.03	1.438	27.72	26.3	5.69	1750	2.118	34.551	0.98	47.7	1.6114					
3000	3001	1.61	34.659	2.20	7.587	2.472	210	2.86	39.5	0.00	0.08	1.384	27.73	28.5	5.53	2000	1.936	34.580	1.23	43.3	1.7456					
3250	3250	1.59	34.667	2.25	7.598	2.473	211	2.86	39.1	0.00	0.08	1.339	27.74	29.5	5.45	2250	1.765	34.624	1.53	39.7	1.8704					
3500	3500	1.58	34.674	2.55	7.611	2.492	209	2.83	38.9	0.00	0.08	1.303	27.74	33.0	5.48	2500	1.702	34.644	1.79	37.5	1.9880					
3750	3749	1.57	34.678	2.64	7.624	2.476	205	2.77	38.1	0.00	0.09	1.266	27.75	34.1	5.10	2750	1.638	34.659	2.02	35.9	2.1039					
																3000	1.601	34.670	2.20	34.9	2.2168					
																3250	1.583	34.677	2.35	34.2	2.3289					
																3500	1.575	34.663	2.50	33.7	2.4415					
																3750	1.449	34.666	3.52	31.8	2.5555					

Table 6. Summary of hydrographic data at Station 1

Station	1	Date	June 17, 1986	Lat.	57° 59.5' N	Air T.	6.4 C	Weather	Fog	Sea	3
Depth	3640 m	TIDE	15:16 - 16:21	Long.	176° 26.5' W	Barro.	999.0 mb	Wind	ENE - 8.5 m/s	Surf	2
CTD data											
RHS of HYDROCAST		D	T	S	D.O.	pH	AT	Sal	DO	Temp	Depth
Sample No.	m	m	°C	m	ml/l	ml/l	mg/l	‰	‰	°C	m
0	0	6.10	33.066	7.72	7.998	2.281	62.3	1.44	14.8	0.15	0.30
10	10	4.46	33.450	7.68	7.929	2.281	41.0	1.73	10.8	0.19	0.61
20	20	3.60	33.236	7.73	7.872	2.293	42.1	1.98	22.7	0.19	0.17
30	30	3.34	33.279	7.71	7.856	2.288	42.9	2.06	23.3	0.22	0.24
40	40	3.20	33.229	7.62	7.843	2.297	43.9	2.07	23.7	0.24	0.37
50	49	2.92	33.233	7.56	7.828	2.303	45.8	2.16	24.4	0.27	0.38
75	76	2.24	33.248	7.34	7.791	2.300	48.0	2.21	25.6	0.33	0.22
100	100	2.13	33.239	7.47	7.798	2.297	48.0	2.19	25.5	0.41	0.15
200	200	3.71	33.587	3.63	7.569	2.322	76.0	2.63	34.1	0.00	0.03
300	300	3.71	33.818	1.07	7.455	2.341	93.3	2.86	39.1	0.00	0.04
500	500	3.44	34.082	0.72	7.399	2.377	115	3.22	42.0	0.00	0.01
750	750	3.09	34.266	0.44	7.386	2.409	136	3.22	42.5	0.00	0.02
1000	1000	2.77	34.379	0.45	7.417	2.433	150	3.20	42.1	0.00	0.03
CTD data											
		D	T	S	DO	Temp	‰	DO	Temp	‰	Depth
		m	°C	m	ml/l	mg/l	%	ml/l	°C	m	m
		0	6.200	32.654	7.69	245.6	0.0000	0	6.200	32.654	7.69
		10	5.936	33.092	7.62	194.7	0.0207	10	5.936	33.092	7.62
		20	3.856	33.191	7.65	165.4	0.0387	20	3.856	33.191	7.65
		30	3.508	33.239	7.37	158.6	0.0552	30	3.508	33.239	7.37
		50	3.280	33.252	7.18	155.4	0.0868	50	3.280	33.252	7.18
		100	2.182	33.250	6.83	146.9	0.1627	100	2.182	33.250	6.83
		150	2.203	33.277	6.68	145.0	0.2349	150	2.203	33.277	6.68
		200	3.692	33.595	4.20	133.3	0.3058	200	3.692	33.595	4.20
		300	3.706	33.840	2.00	144.9	0.4311	300	3.706	33.840	2.00
		800	3.569	33.398	1.18	103.0	0.5126	800	3.569	33.398	1.18
		1000	3.026	34.258	0.40	74.3	0.9055	1000	3.026	34.258	0.40
		1000	3.005	34.327	0.52	71.9	1.0591	1000	3.005	34.327	0.52

Table 7. Summary of hydrographic data at Station E

Station E	Depth	Date		TIME		Lat.		56 - 01.3 N		Air T.		8.4 C		Weather		Drizzle		Sea		4			
		3910 m	T	S	0.0	pH	AT	SiO ₂	Po4	NO ₃	NO ₂	NH ₄	Pot-T	%DO	AOU	D	T	S	O.O.	Dst	De-I		
RMS of HYDROCAST		No.	m	C	m/l	mEq/l	Um	Um	Um	Um	Um	C	m/l	m	C	m/l	m	Um	Um	Um	Um		
0	0	8.40	32.600	7.08	7.964	----	16.7	1.14	11.8	0.12	0.15	0.400	25.33	106.5	-0.43	0	8.239	32.586	7.30	262.0	0.0000		
50	50	6.39	32.661	7.26	7.943	----	20.0	1.23	13.4	0.14	0.26	0.366	25.66	104.3	-0.30	10	8.217	32.612	7.61	259.8	0.0266		
100	100	4.35	33.030	5.86	7.780	----	34.4	1.69	23.3	0.05	0.04	4.343	26.18	80.3	1.43	20	8.158	32.608	7.68	259.3	0.0523		
200	200	3.78	33.913	0.73	7.370	----	90.0	3.22	45.2	0.00	0.01	3.766	26.94	9.9	6.62	30	6.973	32.625	7.75	242.2	0.0772		
300	300	3.72	34.032	0.50	7.375	----	102	3.26	45.5	0.00	0.01	3.699	27.04	6.8	6.86	50	6.284	32.624	7.52	233.9	0.1256		
400	400	3.62	34.115	0.45	7.392	----	113	3.18	45.5	0.00	0.02	3.592	27.12	6.1	6.92	75	4.974	32.678	7.28	215.1	0.1824		
500	500	3.50	34.177	0.39	7.402	----	120	3.20	45.0	0.00	0.01	3.465	27.18	5.3	7.00	100	4.176	32.669	5.50	162.5	0.2295		
600	600	3.38	34.241	0.53	7.411	----	126	3.23	44.8	0.00	0.00	3.338	27.24	7.2	6.88	150	3.820	33.626	2.09	117.1	0.2998		
700	700	3.22	34.295	0.40	7.445	----	136	3.23	45.2	0.00	0.02	3.172	27.30	5.4	7.04	200	3.787	33.925	4.13	109.3	0.3572		
800	800	3.07	34.334	0.43	7.420	----	140	3.02	44.9	0.00	0.01	3.015	27.35	5.8	7.03	300	3.725	34.037	0.66	100.3	0.4638		
900	900	2.92	34.369	0.37	7.426	----	144	3.19	44.8	0.00	0.02	2.858	27.39	4.9	7.12	400	3.598	34.134	0.58	91.7	0.5626		
1000	1000	2.78	34.406	0.47	7.431	----	152	3.19	45.0	0.00	0.01	2.712	27.43	6.3	7.05	500	3.440	34.224	0.54	84.2	0.6540		
1250	1249	2.46	34.475	0.58	7.447	----	162	3.13	44.6	0.00	0.01	2.376	27.51	7.8	6.98	600	3.300	34.276	0.50	78.3	0.7933		
1500	1499	2.21	34.532	0.84	7.473	----	168	3.08	44.0	0.00	0.01	2.109	27.58	11.0	6.78	700	3.164	34.319	0.51	73.8	0.8200		
1750	1750	2.05	34.568	1.16	7.496	----	174	3.11	43.6	0.00	0.01	1.930	27.62	15.2	6.48	800	3.009	34.359	0.52	69.5	0.8964		
2000	2000	1.87	34.604	1.52	7.532	----	177	2.93	42.4	0.00	0.02	1.732	27.66	19.8	6.16	1000	2.749	34.426	0.52	62.2	1.0395		
2250	2249	1.75	34.630	1.82	7.562	----	177	2.88	41.4	0.00	0.01	1.592	27.69	23.6	5.68	1250	2.446	34.491	0.66	54.8	1.2015		
2500	2500	1.68	34.645	2.07	7.587	----	178	2.89	41.0	0.00	0.01	1.501	27.71	26.8	5.65	1500	2.197	34.547	0.88	48.6	1.3480		
2750	2750	1.61	34.659	2.35	7.608	----	178	2.85	40.4	0.00	0.02	1.408	27.73	30.4	5.38	1750	2.054	34.579	1.13	45.0	1.4635		
3000	2999	1.55	34.670	2.63	7.632	----	177	2.70	39.3	0.00	0.02	1.325	27.74	34.0	5.11	2000	1.669	34.617	1.52	40.8	1.6110		
3250	3249	1.50	34.681	2.87	7.653	----	175	2.63	38.8	0.00	0.02	1.251	27.75	37.0	4.88	2250	1.754	34.640	1.86	38.2	1.7930		
3500	3500	1.46	34.690	3.15	7.674	----	171	2.65	37.9	0.00	0.02	1.166	27.76	40.6	4.61	2500	1.676	34.657	2.14	36.4	1.8450		
3750	3750	1.45	34.696	3.25	7.689	----	168	2.64	37.1	0.00	0.02	1.149	27.77	41.9	4.51	2750	1.609	34.671	2.43	34.9	1.9570		
																				33.5	2.0663		
																				32.4	2.1737		
																				3500	1.460	34.702	3.24
																				3750	1.446	34.707	3.44
																					31.0	2.3846	

Table 8. Summary of hydrographic data at Station F

Station F Depth 4110 m	Date		June 29, 1986		Lat.		53 - 02.5 N		Air T.		9.3 C		Weather		Fog		Sea Swell		3					
	TIME	10 : 48 - 14 : 04	Long.	145 - 00.0 W	Baro.	1008.1 mb	Wind	SE - 7.0 m/s	D.O.	m/l	S	m/l	Det	Dej-D	CTD0 data	D	T	%	AOU	m/l	C	m/l	C	m/l/t
<u>AMS or HYDROCAST</u>																								
Sample No.	T	S	O.O.	pH	AT	SiO ₂	Po4	N03	N02	NH4	PoC-T	Sig-T	%DO	AOU	D	T	S	D.O.	m/l	C	m/l	C	m/l/t	
No.	m	m	ml/l	ml/l	deg	um	um	um	um	um	um	um	%	ml/l	m	m	%	ml/l	m	m	m	m	m	
0	0	7.98	32.400	7.21	7.952	2.246	14.4	1.45	11.2	0.15	0.13	7.980	25.24	-0.49	0	7.981	32.294	5.47	200.3	0.0000				
50	50	6.13	32.543	7.38	7.935	2.257	16.6	1.23	11.6	0.16	0.21	6.126	25.60	-0.37	10	7.988	32.488	5.42	254.5	0.0281				
100	100	4.24	32.301	6.60	7.930	2.249	27.1	1.65	19.7	0.20	0.03	4.233	26.08	90.2	0.72	20	7.752	32.497	5.52	222.0	0.0548			
200	200	3.65	33.845	1.46	7.408	2.334	84.1	3.02	42.9	0.00	0.02	3.836	26.90	19.6	5.92	30	6.913	32.531	5.63	247.3	0.0803			
300	299	3.68	33.980	0.69	7.370	2.339	98.5	3.19	44.0	0.00	0.03	3.658	27.01	9.4	6.68	50	6.290	32.546	5.58	239.6	0.1285			
400	399	3.62	34.091	0.67	7.400	2.353	111	3.12	45.0	0.00	0.04	3.592	27.10	9.1	6.70	75	5.569	32.549	5.52	231.2	0.1875			
500	500	3.53	34.164	0.62	7.409	2.366	110	3.11	45.0	0.00	0.03	3.495	27.17	8.4	6.77	100	4.085	33.128	4.76	112.3	0.2399			
600	600	3.40	34.233	0.59	7.420	2.380	126	---	44.6	0.00	0.01	3.358	27.24	8.0	6.82	150	3.599	33.719	2.32	123.1	0.3127			
700	701	3.25	34.292	0.55	7.423	2.388	134	3.13	44.6	0.00	0.02	3.201	27.30	7.4	6.90	200	3.620	33.970	1.38	111.9	0.3725			
800	799	3.09	34.337	0.51	7.426	2.404	139	3.17	44.8	0.00	0.03	3.035	27.35	6.8	6.95	300	3.679	34.009	0.65	101.9	0.4815			
900	900	2.93	34.375	0.52	7.429	2.405	144	3.16	44.5	0.00	0.02	2.868	27.39	6.9	6.97	400	3.616	34.104	0.60	94.1	0.5815			
1000	1000	2.79	34.407	0.47	7.431	2.412	152	3.15	45.2	0.00	0.04	2.722	27.43	6.3	7.04	500	3.502	34.187	0.60	86.9	0.6754			
1250	1250	2.49	34.472	0.60	7.444	2.426	163	3.15	45.0	0.00	0.02	2.405	27.51	7.9	6.97	600	3.376	34.257	0.64	80.4	0.7634			
1500	1500	2.26	34.524	0.81	7.465	2.441	167	3.09	43.8	0.00	0.01	2.158	27.57	10.6	6.80	700	3.251	34.306	0.69	75.6	0.8461			
1750	1750	2.07	34.565	1.12	7.492	2.443	174	3.01	43.9	0.00	0.04	1.950	27.62	14.7	6.52	800	3.071	34.356	0.58	70.3	0.9241			
2000	2000	1.94	34.596	1.44	7.519	2.454	175	2.98	43.2	0.00	0.04	1.801	27.65	10.8	6.23	1000	2.784	34.421	0.52	62.9	1.0689			
2250	2249	1.81	34.621	1.75	7.547	2.454	176	2.91	41.6	0.00	0.01	1.651	27.68	22.8	5.94	1250	2.483	34.486	0.57	55.5	1.2328			
2500	2499	1.71	34.640	2.06	7.575	2.458	177	2.84	41.3	0.00	0.01	1.530	27.71	26.7	5.65	1500	2.240	34.539	0.76	49.5	1.3815			
2750	2750	1.63	34.659	2.37	7.601	2.466	176	2.78	40.6	0.00	0.01	1.428	27.73	30.7	5.36	1750	2.067	34.578	1.04	45.2	1.5188			
3000	3001	1.57	34.670	2.64	7.626	2.472	174	2.74	38.9	0.00	0.00	1.345	27.74	34.1	5.10	2000	1.921	34.610	1.33	41.7	1.6475			
3250	3251	1.52	34.680	2.89	7.646	2.458	174	2.67	38.4	0.00	0.04	1.271	27.75	37.3	4.86	2250	1.803	34.634	1.63	39.0	1.7696			
3500	3500	1.49	34.688	3.10	7.664	2.468	173	2.61	38.4	0.00	0.01	1.215	27.76	40.0	4.65	2500	1.707	34.654	1.95	36.9	1.8866			
3750	3750	1.47	34.694	3.30	7.676	2.461	169	2.53	37.0	0.00	0.12	1.169	27.77	42.6	4.46	2750	1.630	34.669	2.25	35.2	1.9999			
4000	4000	1.47	34.698	3.38	7.683	2.467	170	2.55	37.5	0.00	0.03	1.141	27.77	43.6	4.38	3000	1.571	34.681	2.52	33.8	2.1105			
																3250	1.525	34.691	2.80	32.7	2.2189			
																3500	1.488	34.699	3.03	31.8	2.3259			
																4000	1.473	34.709	3.36	31.0	2.5403			

Table 9. Summary of hydrographic data at Station G

Station	6	Date	June 30, 1966			Lat.	50 - 00.5 N			Air T.	7.7 C			Weather	Overcast	Sea	3					
			Depth	4260 m	TIME		Long.	144	- 59.2 W		Baro.	998.9 mb	Wind	SE -10.0 m/s	Swell	3						
CTD0 data																						
AMS or HYDROCAST	Sample	D	T	S	D.O.	pH	AT	S102	P04	N03	N02	NH4	Pot-T	Sig-t	%D0	AOU	D	T	S	D.O.	Dist	De1-D
No.	m	C	m/l	m/l	mg/l	UH	UH	UH	UH	UH	UH	UH	UH	UH	m/l	m	C	m/l	m/l	m/l	m/l/t	
0	0	9.13	32.532	7.08	7.980	-----	9.5	4.03	8.60	0.09	0.10	9.130	25.17	108.2	-0.54	0	9.125	32.550	6.57	277.6	0.0000	
50	50	7.81	32.576	7.23	7.969	-----	10.0	4.10	9.90	0.10	0.10	7.805	25.40	107.3	-0.49	10	9.123	32.540	6.59	278.3	0.0279	
100	99	6.23	32.749	6.89	7.922	-----	12.8	4.25	12.7	0.34	0.10	6.221	25.75	98.6	0.09	20	9.086	32.539	6.64	277.9	0.0574	
200	200	4.86	33.821	3.27	7.603	-----	55.2	2.40	33.2	0.00	0.06	4.844	26.76	45.6	3.90	30	9.048	32.543	6.95	276.9	0.0843	
300	300	4.42	33.972	1.84	7.485	-----	75.9	0.00	0.02	4.397	26.92	25.4	5.40	50	8.217	32.576	6.92	262.4	0.1384			
400	400	4.09	34.020	1.28	7.447	-----	91.5	3.01	42.0	0.00	0.02	4.061	27.00	17.6	6.01	75	7.596	32.574	6.69	254.2	0.2044	
500	500	3.87	34.108	0.94	7.435	-----	104	3.07	41.8	0.00	0.02	3.634	27.03	42.8	6.39	100	6.251	32.669	6.60	230.1	0.2646	
600	600	3.66	34.163	0.79	7.431	-----	115	3.09	43.4	0.00	0.01	3.617	27.17	10.7	6.57	150	5.137	33.695	4.23	140.3	0.3671	
700	700	3.49	34.241	0.67	7.425	-----	124	3.12	43.9	0.00	0.02	3.440	27.23	9.1	6.72	200	4.812	33.846	3.05	125.5	0.4252	
800	802	3.30	34.288	0.50	7.422	-----	130	3.18	44.4	0.00	0.01	3.243	27.29	6.7	6.92	300	4.330	33.952	1.85	112.5	0.5462	
900	900	3.12	34.336	0.47	7.422	-----	141	3.18	44.7	0.00	0.00	3.057	27.34	6.3	6.99	400	4.049	34.052	4.34	102.2	0.6571	
1000	999	2.94	34.376	0.45	7.424	-----	147	3.19	45.2	0.00	0.00	2.871	27.38	6.0	7.04	500	3.843	34.128	4.19	94.5	0.7584	
1250	1250	2.61	34.450	0.46	7.433	-----	154	3.19	43.8	0.00	0.00	2.584	27.48	6.1	7.09	600	3.674	34.197	0.94	87.7	0.8542	
1500	1500	2.35	34.504	0.72	7.453	-----	165	3.14	44.7	0.00	0.00	2.247	27.55	9.5	6.87	700	3.479	34.258	0.70	81.3	0.9437	
1750	1749	2.13	34.555	1.07	7.483	-----	172	3.07	43.8	0.00	0.00	2.009	27.60	14.0	6.56	800	3.274	34.312	0.60	75.4	1.0276	
2000	2000	1.97	34.589	1.34	7.512	-----	176	2.98	42.9	0.00	0.00	1.630	27.64	17.5	6.32	1000	2.981	34.387	0.49	66.9	1.1820	
2250	2243	1.84	34.616	1.68	7.540	-----	176	2.91	42.0	0.00	0.00	1.681	27.68	21.6	6.03	1250	2.603	34.465	0.50	58.0	1.3546	
2500	2500	1.73	34.637	2.04	7.571	-----	176	2.82	41.3	0.00	0.00	1.550	27.70	26.5	5.67	1500	2.330	34.522	0.74	51.5	1.5104	
2750	2749	1.65	34.663	2.32	7.598	-----	174	2.77	39.7	0.00	0.00	1.448	27.72	30.0	5.40	1750	2.106	34.571	4.04	46.1	1.6515	
3000	3000	1.58	34.667	2.63	7.622	-----	174	2.74	39.5	0.00	0.00	1.355	27.74	34.0	5.14	2000	1.951	34.603	1.36	42.5	1.7825	
3250	3251	1.53	34.679	2.88	7.643	-----	173	2.66	38.8	0.00	0.00	1.280	27.75	37.2	4.86	2250	1.829	34.629	4.67	39.6	1.9068	
3500	3500	1.50	34.686	3.06	7.659	-----	171	2.61	38.4	0.00	0.00	1.225	27.76	39.5	4.69	2500	1.714	34.651	2.03	37.1	2.0250	
3750	3749	1.49	34.691	3.23	7.672	-----	171	2.60	37.9	0.00	0.00	1.188	27.76	41.7	4.52	2750	1.640	34.666	2.34	35.4	2.1387	
4000	4000	1.49	34.696	3.32	7.683	-----	171	2.55	37.2	0.00	0.00	1.160	27.77	42.8	4.43	3000	1.580	34.678	2.63	34.4	2.2499	
																	3250	1.532	34.669	2.92	33.0	2.3594
																	3500	1.592	34.697	3.45	32.1	2.4672
																	3750	1.466	34.703	3.34	31.6	2.5747

Table 10. Summary of hydrographic data at Station F'

Station	F'	Date	July 14/15, 1986			Lat.			52 - 59.9 N			Air T.			8.7 C			Weather			Rain			
			Depth	4090 m	TIME	22:27	- 01:56	Long.	144	- 155.9 W	Barto.	1028.4 mb	Wind	S - 5.5 m/s	Swell	3	3	3	3	3	3	3	3	3
No.	Sample	D	T	S	D.O.	pH	AT	SiO ₂	PO ₄	NO ₃	NH ₄	Ptot-T	Sig-t	%DD	ADU	CTDO data			CDO data			CDO data		
																ml/l	m	UN	UN	UN	UN	UN	UN	UN
0	0	8.73	32.446	7.05	7.961	2.247	13.9	1.14	10.1	0.07	0.26	0.730	25.16	106.7	-0.44	0	8.729	32.455	6.75	278.8	0.0000			
50	50	6.33	32.529	7.49	7.934	2.265	15.0	1.17	10.9	0.06	0.18	0.326	25.56	107.3	-0.51	10	6.737	32.455	6.83	278.9	0.0300			
100	100	4.75	32.503	6.88	7.931	2.266	25.5	1.61	18.6	0.07	0.06	4.743	25.96	92.3	0.55	20	6.635	32.455	6.84	277.4	0.0564			
200	200	3.61	33.836	1.48	7.405	2.336	82.2	3.10	41.9	0.00	0.04	3.597	26.90	20.0	5.91	30	6.540	32.452	6.81	276.3	0.0847			
300	300	3.69	33.984	0.68	7.363	2.365	97.2	3.27	45.2	0.00	0.04	3.668	27.01	9.2	6.69	50	6.174	32.531	6.98	239.5	0.1356			
400	401	3.62	34.084	0.61	7.368	2.372	108	3.24	44.7	0.00	0.07	3.592	27.10	0.3	6.77	75	5.031	32.605	6.82	221.2	0.1933			
500	500	3.49	34.167	0.61	7.400	2.386	117	3.18	44.3	0.00	0.04	3.455	27.17	0.2	6.78	100	4.360	32.809	6.35	199.0	0.2464			
600	599	3.33	34.236	0.53	7.410	2.402	125	3.17	44.4	0.00	0.03	3.289	27.24	7.4	6.89	150	3.657	33.719	2.70	123.6	0.3273			
700	700	3.17	34.289	0.59	7.419	2.405	131	3.18	44.3	0.00	0.04	3.122	27.30	7.9	6.86	200	3.594	33.805	1.54	112.0	0.3668			
800	800	3.02	34.334	0.53	7.423	2.414	138	3.16	44.3	0.00	0.04	2.965	27.35	7.1	6.94	300	3.666	34.026	0.65	100.5	0.4953			
900	900	2.89	34.373	0.61	7.439	2.427	142	3.19	44.5	0.00	0.00	2.829	27.39	8.1	6.88	400	3.508	34.126	0.64	92.3	0.5939			
1000	1001	2.78	34.401	0.49	7.432	2.418	146	3.17	44.6	0.00	0.03	2.712	27.43	6.5	7.03	500	3.449	34.198	0.60	85.5	0.6865			
1250	1250	2.50	34.467	0.59	7.439	2.455	156	3.17	44.5	0.00	0.01	2.415	27.50	7.8	6.98	600	3.294	34.260	0.58	79.5	0.7732			
1500	1500	2.29	34.513	0.77	7.460	2.453	163	3.15	44.4	0.00	0.01	2.188	27.56	10.1	6.83	700	3.145	34.308	0.60	74.5	0.8549			
1750	1749	2.09	34.558	1.15	7.487	2.459	168	3.03	43.8	0.00	0.05	1.970	27.61	15.1	6.49	800	3.023	34.345	0.58	70.7	0.9323			
2000	2000	1.93	34.590	1.42	7.523	2.466	170	2.98	42.3	0.00	0.03	1.791	27.65	19.5	6.25	1000	2.767	34.417	0.55	63.0	1.0772			
2250	2250	1.81	34.617	1.73	7.548	2.468	171	2.89	41.9	0.00	0.04	1.651	27.68	22.5	5.96	1250	2.512	34.476	0.58	56.5	1.2424			
2500	2500	1.71	34.637	2.11	7.575	2.481	173	2.81	41.1	0.00	0.04	1.530	27.70	27.4	5.60	1500	2.289	34.526	0.78	50.9	1.3945			
2750	2750	1.63	34.653	2.41	7.601	2.480	171	2.91	40.0	0.00	0.04	1.428	27.72	31.2	5.32	1750	2.092	34.569	1.07	46.1	1.5351			
3000	2999	1.57	34.666	2.71	7.625	2.480	170	2.70	39.5	0.00	0.03	1.345	27.74	35.0	5.03	2000	1.930	34.604	1.42	42.3	1.6658			
3250	3250	1.52	34.676	2.92	7.647	2.475	169	2.66	38.8	0.00	0.04	1.271	27.75	37.7	4.83	2250	1.796	34.631	1.78	39.2	1.7887			
3500	3498	1.49	34.683	3.08	7.661	2.478	167	2.63	38.0	0.00	0.03	1.215	27.76	39.7	4.87	2500	1.699	34.650	2.10	37.0	1.9061			
3750	3750	1.47	34.688	3.33	7.677	2.471	166	2.57	37.8	0.00	0.24	1.169	27.76	42.9	4.43	2750	1.632	34.664	2.40	35.5	2.0399			
4000	4000	1.47	34.688	3.42	7.693	2.482	164	2.62	37.2	0.00	0.04	1.141	27.76	44.1	4.34	3000	1.565	34.677	2.73	34.1	2.1310			
																3250	1.517	34.687	3.00	33.0	2.2402			
																4000	1.470	34.704	3.59	31.3	2.5636			

Table 11. Summary of hydrographic data at Station C'

Station	C'			Date July 22, 1986			Lat.			53 - 30.5 N			Air T.			7.5 C			Weather Rain			Sea 2		
	Depth	3930 m	TIME	12:21 - 15:11	Long.	177 - 31.7 E	Barro.	1010.8 mb	Wind	E-5.0 m/s	Swell	1	Barro.	1010.8 mb	Wind	E-5.0 m/s	Swell	1	Barro.	1010.8 mb	Wind	E-5.0 m/s	Swell	1
PHS or HYDROCAST												CTD data												
No.	m	C	T	S	D.O.	pH	AT	SiO ₂	PO ₄	NO ₃	NO ₂	NH ₄	Flu-T	Sig-T	%DO	AOU	D	I	S	D.O.	Dst	Dst	Dst	Dst
Sample No.	m	C	T	S	D.O.	pH	AT	SiO ₂	PO ₄	NO ₃	NO ₂	NH ₄	Flu-T	Sig-T	%DO	AOU	D	I	S	D.O.	Dst	Dst	Dst	Dst
0	0	7.55	32.722	7.11	7.910	2.266	17.6	1.57	15.8	0.49	0.46	7.550	25.55	105.0	-0.34	0	7.553	32.741	7.07	241.1	0.0000			
50	50	6.00	33.023	7.14	7.881	2.297	20.1	1.64	17.4	0.26	0.56	5.986	25.99	101.9	-0.13	10	7.528	32.802	6.58	256.2	0.0244			
100	100	3.30	33.207	6.94	7.775	2.293	42.7	2.08	26.0	0.24	0.07	3.234	26.43	91.5	0.54	20	7.354	32.859	6.76	259.7	0.0476			
200	200	3.84	33.667	2.85	7.497	2.324	72.9	2.79	37.6	0.00	0.06	3.826	26.74	38.7	4.51	30	7.176	32.901	6.84	224.2	0.0708			
300	300	3.80	33.911	1.35	7.402	2.353	90.5	3.07	42.1	0.00	0.04	3.779	26.94	18.4	6.00	50	6.218	32.984	6.78	205.3	0.1142			
400	400	3.65	34.019	0.86	7.381	2.370	104	3.16	44.6	0.00	0.04	3.622	27.04	11.7	6.51	75	3.837	33.417	6.73	170.8	0.1619			
500	500	3.52	34.117	0.65	7.382	2.381	116	3.16	45.2	0.00	0.04	3.495	27.13	8.8	6.74	100	3.339	33.483	6.17	161.3	0.2023			
600	600	3.42	34.176	0.63	7.390	2.389	123	3.21	45.1	0.00	0.04	3.378	27.19	8.5	6.78	150	3.523	33.446	4.46	143.0	0.2799			
700	699	3.29	34.227	0.55	7.394	2.402	129	3.17	45.3	0.00	0.03	3.241	27.24	7.4	6.88	200	3.820	33.721	2.80	155.0	0.3472			
800	800	3.15	34.274	0.51	7.397	2.406	134	3.14	45.3	0.00	0.03	3.095	27.29	6.8	6.94	300	3.778	33.932	1.37	108.7	0.4654			
900	900	3.03	34.311	0.49	7.400	2.415	142	3.18	45.4	0.00	0.04	2.968	27.33	6.6	6.98	400	3.646	34.042	0.93	99.2	0.5726			
1000	1000	2.90	34.350	0.52	7.405	2.417	147	3.18	45.4	0.00	0.04	2.831	27.38	6.9	6.98	500	3.537	34.125	0.73	91.9	0.6715			
1250	1250	2.56	34.440	0.56	7.424	2.426	161	3.17	45.1	0.00	0.03	2.475	27.48	7.4	7.00	600	3.443	34.182	0.64	86.7	0.7643			
1500	1500	2.28	34.508	0.76	7.443	2.449	175	3.12	44.6	0.00	0.04	2.178	27.55	10.0	6.85	700	3.301	34.237	0.57	81.3	0.8532			
1750	1750	2.05	34.560	1.03	7.473	2.456	187	3.09	43.2	0.00	0.03	1.990	27.62	13.5	6.62	800	3.145	34.290	0.52	75.9	0.9369			
2000	2001	1.86	34.600	1.35	7.502	2.467	199	3.02	41.8	0.00	0.01	1.722	27.66	17.6	6.39	1000	2.880	34.370	0.49	67.5	1.0918			
2250	2250	1.76	34.624	1.59	7.526	2.465	204	2.94	40.9	0.00	0.01	1.602	27.69	20.6	6.11	1250	2.568	34.452	0.57	58.7	1.2659			
2500	2500	1.68	34.644	1.82	7.547	2.467	209	2.90	40.2	0.00	0.03	1.504	27.71	23.6	5.90	1500	2.279	34.520	0.76	51.3	1.4213			
2750	2750	1.63	34.657	2.02	7.564	2.468	212	2.92	39.5	0.00	0.01	1.428	27.73	26.1	5.71	1750	2.055	34.571	1.01	45.7	1.5617			
3000	3000	1.59	34.666	2.17	7.580	2.470	214	2.82	38.6	0.00	0.03	1.364	27.74	28.1	5.56	2000	1.874	34.610	1.33	41.4	1.6902			
3250	3250	1.58	34.674	2.32	7.592	2.477	215	2.84	39.0	0.00	0.03	1.329	27.74	30.0	5.41	2250	1.774	34.633	1.60	38.9	1.8112			
3500	3500	1.57	34.679	2.36	7.605	2.477	216	2.82	38.8	0.00	0.00	1.253	27.75	30.5	5.38	2500	1.692	34.653	1.63	36.8	1.9278			
3750	3750	1.57	34.684	2.53	7.616	2.482	211	2.72	38.7	0.00	0.04	1.286	27.75	32.7	5.21	2750	1.636	34.667	2.06	35.4	2.0412			
3850	3879	1.57	34.686	2.60	7.627	2.480	206	2.76	37.4	0.00	0.04	1.251	27.75	33.6	5.14	3000	1.597	34.678	2.24	34.2	2.1527			
3500	3500	1.575	34.690	2.54												3250	1.581	34.685	2.42	33.6	2.2633			
3750	3750	1.575	34.694	2.70												3500	1.575	34.690	2.54	33.4	2.3745			

Table 12. Summary of hydrographic data at Station H

Station No.	H	Date			July 25, 1986			Lat.			51° 03' 6 N			Air T.			9.6 C			Weather			Rain			Sea				
		Depth	4760 m	TIME	00 : 22 - 03 : 32	Long.	170 - 58.1 E	Baro.	1011.4 mb	Wind	W -7.0 m/s	Swell	1	CDDO data	0	T	S	D.0.	Dst	Del-D										
RHS OF HYDROCAST		D	T	S	0.0.	pH	AT	Sig/I	P04	N03	N02	NH4	Pot-T	Sig-t	%-D0	AOU	D	T	S	ml/l	ml/l	ml/l	ml/l	ml/l	ml/l	ml/l	ml/l	ml/l	ml/l	
0	0	9.00	32.589	7.01	7.991	2.263	16.5	1.12	11.3	0.11	0.25	9.000	25.23	106.9	-0.45	0	9.004	32.597	4.36	272.2	0.0000									
50	49	3.82	32.950	7.11	7.853	2.283	27.1	1.72	20.2	0.50	0.53	3.817	26.17	96.2	0.28	10	8.986	32.599	4.23	272.0	0.0279									
100	100	3.91	33.350	5.23	7.695	2.306	50.7	2.21	29.5	0.00	0.11	3.903	26.48	71.1	2.13	20	8.220	32.718	4.28	252.0	0.0545									
150	200	3.79	33.740	2.51	7.498	2.338	79.0	2.80	39.0	0.00	0.23	3.776	26.81	34.1	4.85	30	7.394	32.688	4.34	243.0	0.0787									
200	300	3.72	33.923	1.38	7.426	2.350	96.7	3.03	43.1	0.00	0.08	3.693	26.96	18.7	5.98	50	4.092	32.682	4.24	190.1	0.1226									
300	400	3.57	34.045	1.00	7.413	2.369	110	3.11	44.0	0.00	0.15	3.542	27.07	13.5	6.39	75	3.736	33.061	3.91	174.1	0.1687									
400	500	3.46	34.131	0.82	7.410	2.376	119	3.15	44.5	0.00	0.05	3.425	27.15	11.1	6.58	100	3.839	35.316	3.33	155.7	0.2091									
500	600	3.37	34.199	0.67	7.411	2.386	126	3.16	44.8	0.00	0.38	3.328	27.21	9.0	6.75	150	4.042	35.593	2.34	136.8	0.2837									
600	700	3.22	34.261	0.52	7.415	2.391	134	3.17	45.0	0.00	0.05	3.172	27.27	7.0	6.92	200	3.886	35.753	1.75	123.2	0.3491									
700	800	3.07	34.307	0.56	7.419	2.402	142	3.19	45.0	0.00	0.08	3.015	27.33	7.5	6.91	300	3.718	33.940	0.98	107.5	0.4669									
800	900	2.96	34.344	0.58	7.424	2.416	146	3.18	45.0	0.00	0.17	2.898	27.37	7.7	6.90	400	3.546	34.080	0.70	96.9	0.5709									
900	1000	2.80	34.383	0.59	7.430	2.423	152	3.16	45.0	0.00	0.05	2.732	27.41	7.9	6.92	500	3.451	34.153	0.57	88.9	0.6672									
1000	1250	2.44	34.472	0.70	7.450	2.439	163	3.11	44.8	0.00	0.03	2.356	27.51	9.2	6.88	600	3.336	34.222	0.52	82.7	0.7573									
1250	1500	2.21	34.526	0.97	7.475	2.447	169	3.08	43.8	0.00	0.02	2.109	27.58	12.7	6.85	700	3.205	34.200	0.46	77.2	0.8418									
1500	1750	2.03	34.566	1.31	7.506	2.453	171	2.98	43.3	0.00	0.00	1.910	27.62	17.1	6.34	800	3.061	34.325	0.45	72.4	0.9218									
1750	2000	1.89	34.599	1.55	7.532	2.478	177	2.91	42.4	0.00	0.05	1.751	27.66	20.2	6.13	1000	2.801	34.395	0.47	65.0	1.0703									
2000	2250	1.76	34.626	1.89	7.563	2.484	178	2.88	41.2	0.00	0.02	1.602	27.69	24.5	5.81	1250	2.462	34.479	0.58	55.9	1.2372									
2250	2500	1.68	34.644	2.25	7.591	2.485	176	2.82	40.5	0.00	0.02	1.501	27.74	29.2	5.47	1500	2.200	34.541	0.81	49.1	1.3853									
2500	2750	1.62	34.660	2.55	7.816	2.487	171	2.74	39.8	0.00	0.02	1.418	27.73	33.0	5.18	1750	2.020	34.581	1.08	44.6	1.5210									
2750	3000	1.55	34.672	2.86	7.643	2.482	167	2.65	38.3	0.00	0.05	1.325	27.74	36.9	4.88	2000	1.884	34.611	1.32	41.4	1.6481									
3000	3250	1.52	34.681	3.01	7.658	2.487	163	2.64	38.3	0.00	0.05	1.271	27.75	38.9	4.74	2250	1.762	34.638	1.59	38.4	1.7669									
3250	3500	1.49	34.686	3.20	7.673	2.495	159	2.57	37.9	0.00	0.00	1.215	27.76	41.3	4.55	2500	1.670	34.657	1.90	36.4	1.8840									
3500	3750	1.46	34.693	3.39	7.684	2.493	157	2.53	37.6	0.00	0.03	1.159	27.77	43.7	4.37	2750	1.612	34.671	2.15	34.9	1.9960									
3750	4000	1.45	34.696	3.48	7.693	2.491	153	2.53	36.9	0.00	—	1.121	27.77	44.8	4.28	3000	1.552	34.692	2.38	33.6	2.1056									
3999	4000	—	—	—	—	—	—	—	—	—	—	—	—	—	—	3250	1.518	34.690	2.56	32.7	2.2135									
3500	4000	—	—	—	—	—	—	—	—	—	—	—	—	—	—	3500	1.482	34.697	2.76	32.9	2.3206									
3999	4000	—	—	—	—	—	—	—	—	—	—	—	—	—	—	4000	1.450	34.706	3.07	31.1	2.5345									

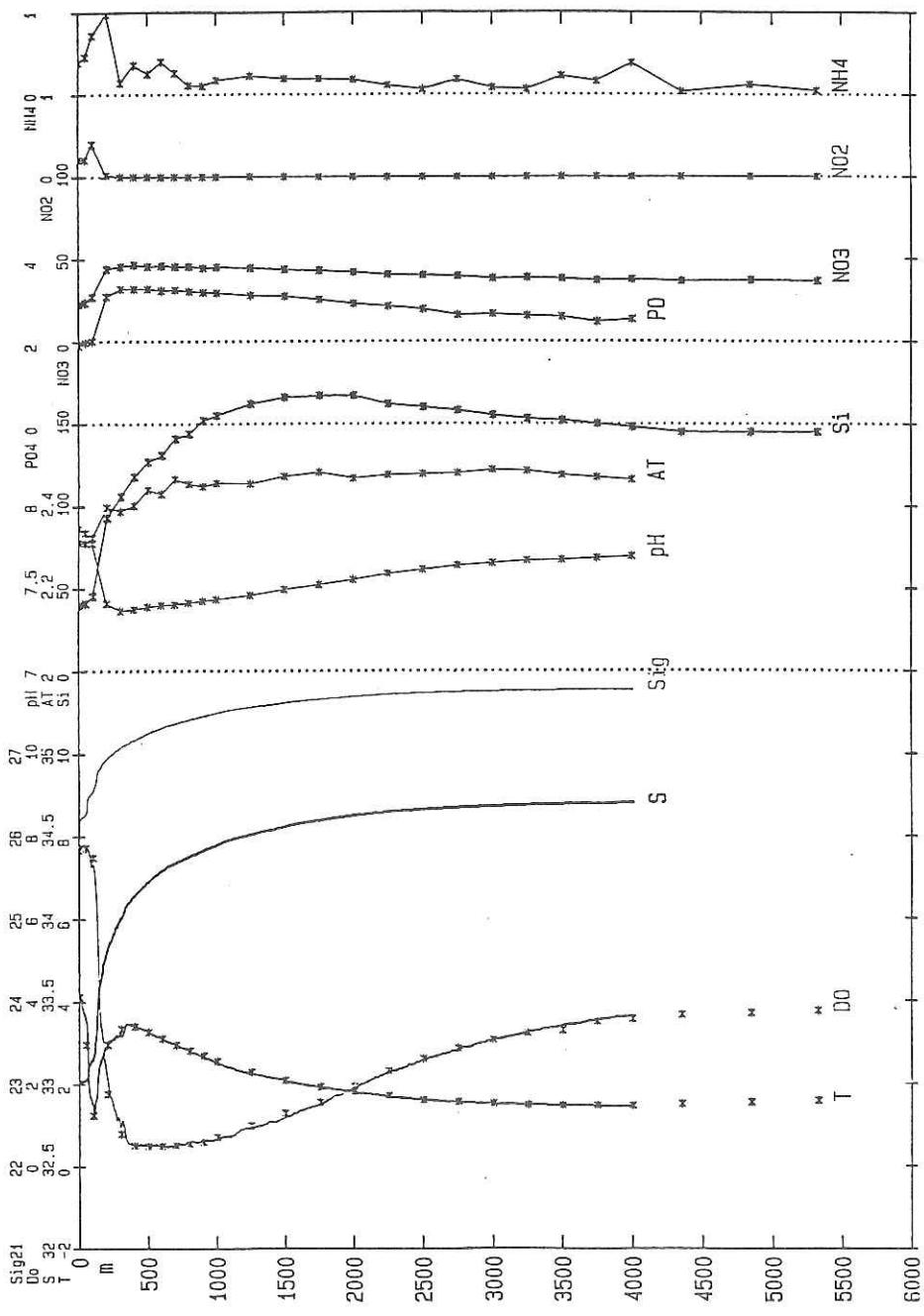


Fig. 2. Vertical profiles of temperature, salinity, sigma-t, dissolved oxygen, pH, alkalinity, silicic acid, phosphate, nitrate, nitrite and ammonium at Station B.

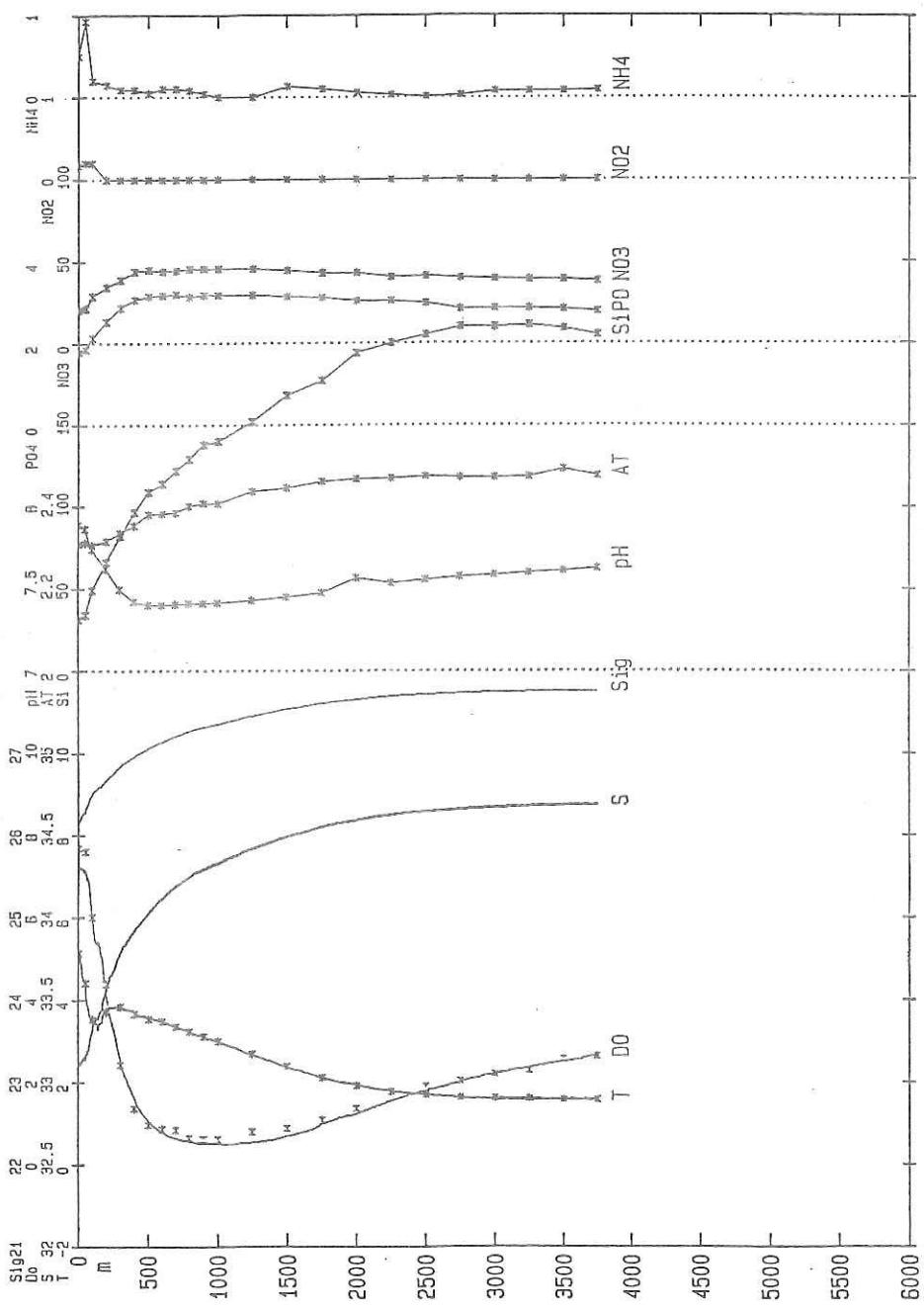


Fig. 3. Vertical profiles of temperature, salinity, sigma-t, dissolved oxygen, pH, alkalinity, silicic acid, phosphate, nitrate, nitrite and ammonium at Station C.

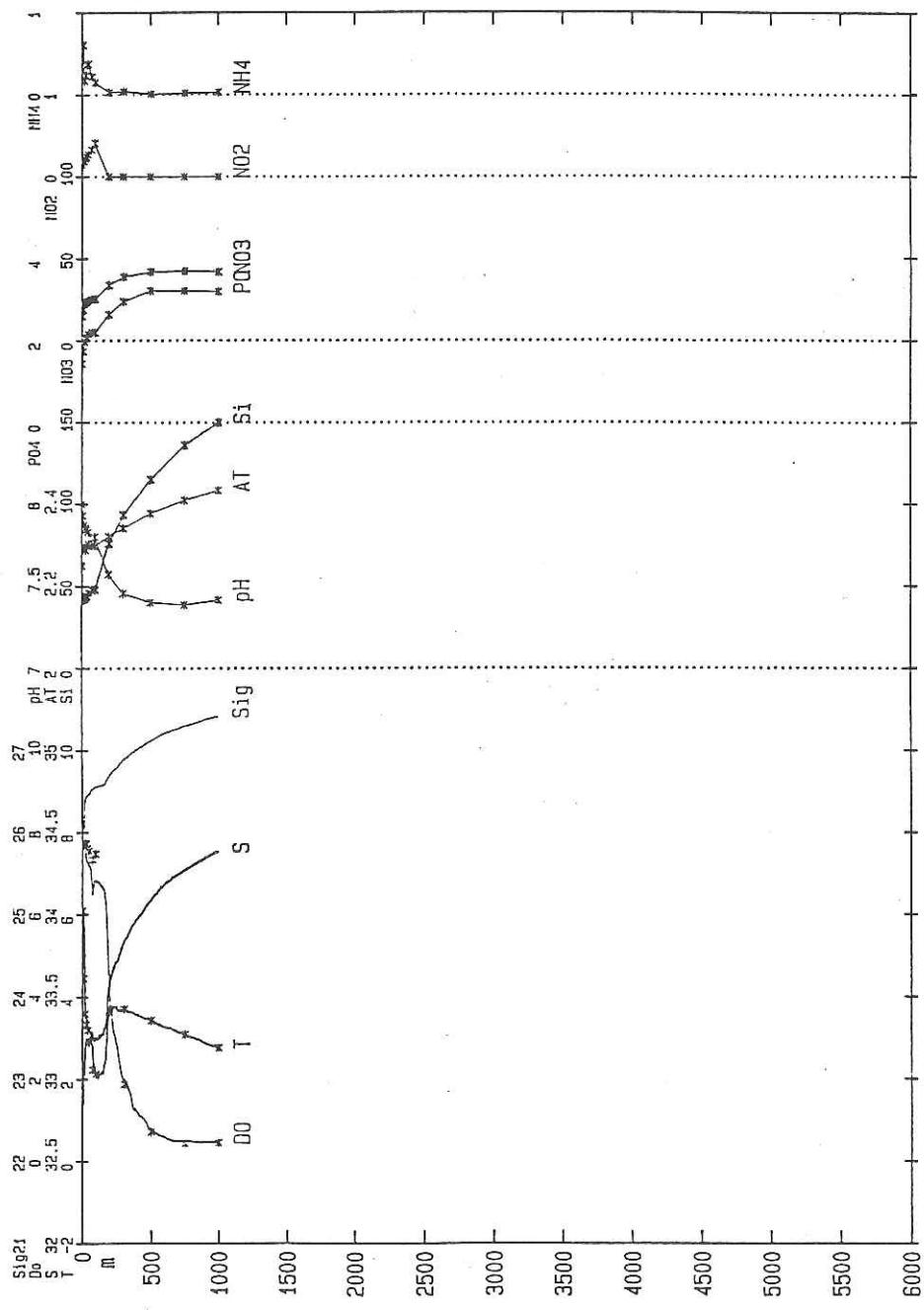


Fig. 4. Vertical profiles of temperature, salinity, sigma-t, dissolved oxygen, pH, alkalinity, silicic acid, phosphate, nitrate, nitrite and ammonium at Station 1.

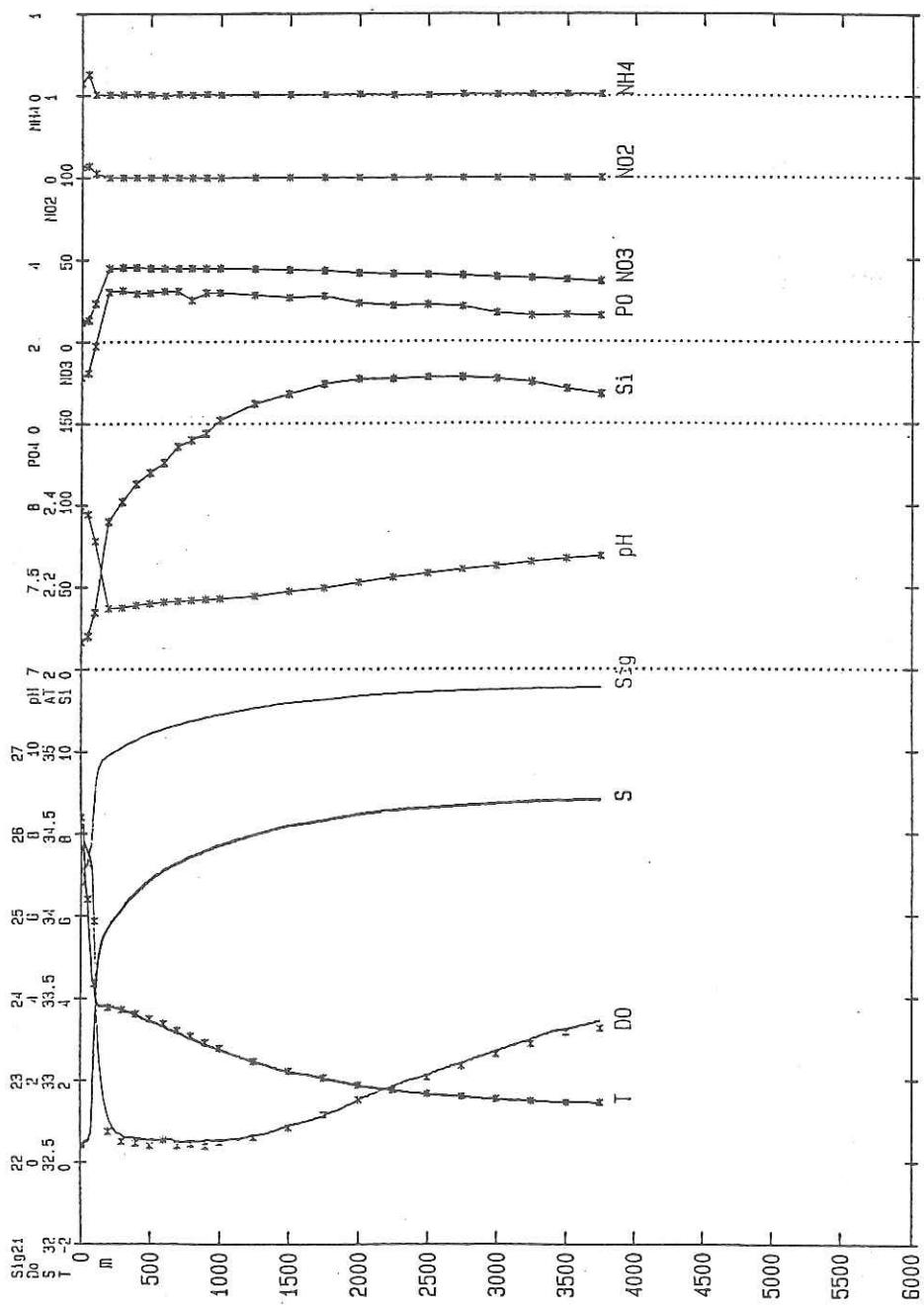


Fig. 5. Vertical profiles of temperature, salinity, sigma-t, dissolved oxygen, pH, alkalinity, silicic acid, phosphate, nitrate, nitrite and ammonium at Station E.

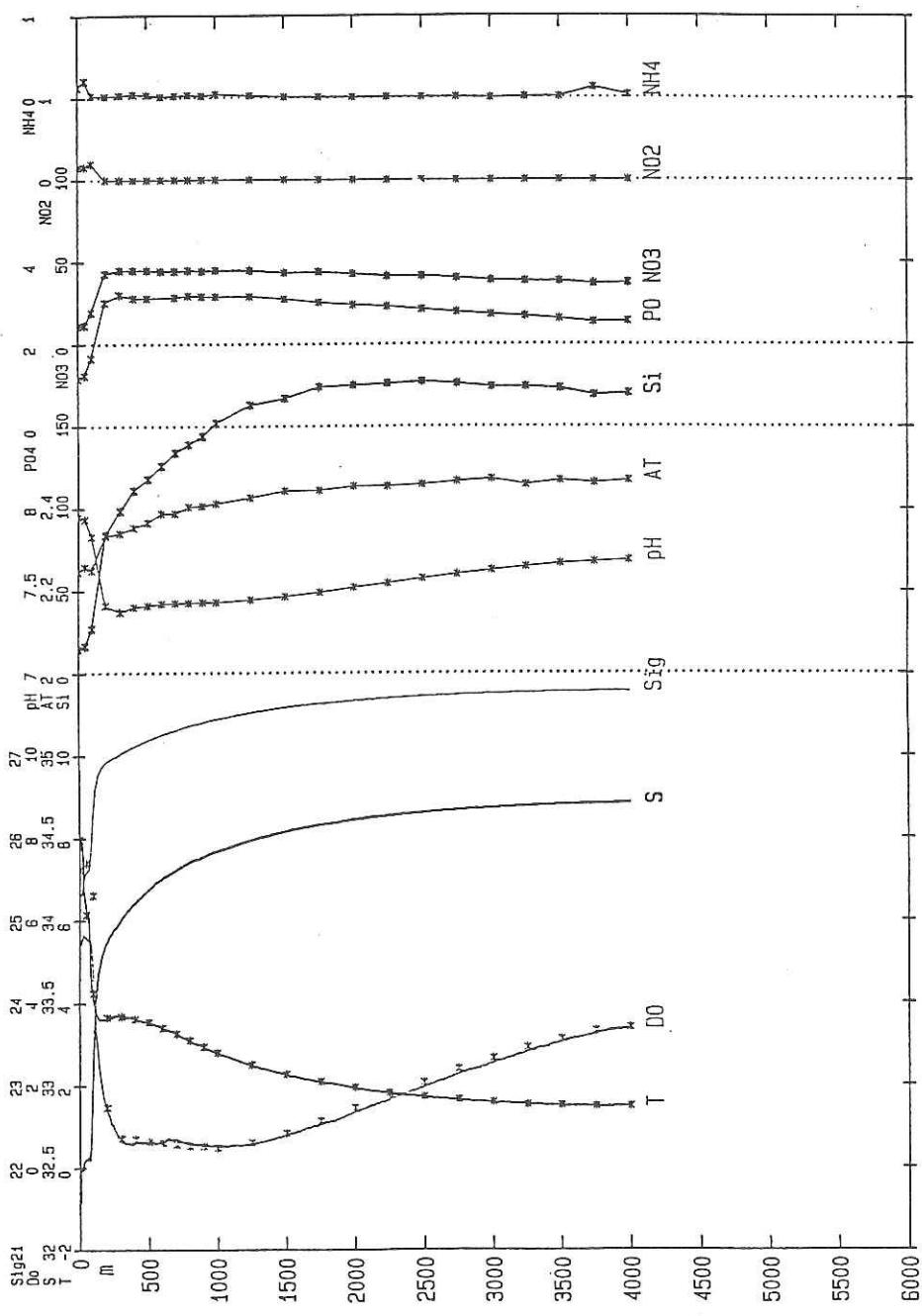


Fig. 6. Vertical profiles of temperature, salinity, sigma-t, dissolved oxygen, pH, alkalinity, silicic acid, phosphate, nitrate, nitrite and ammonium at Station F.

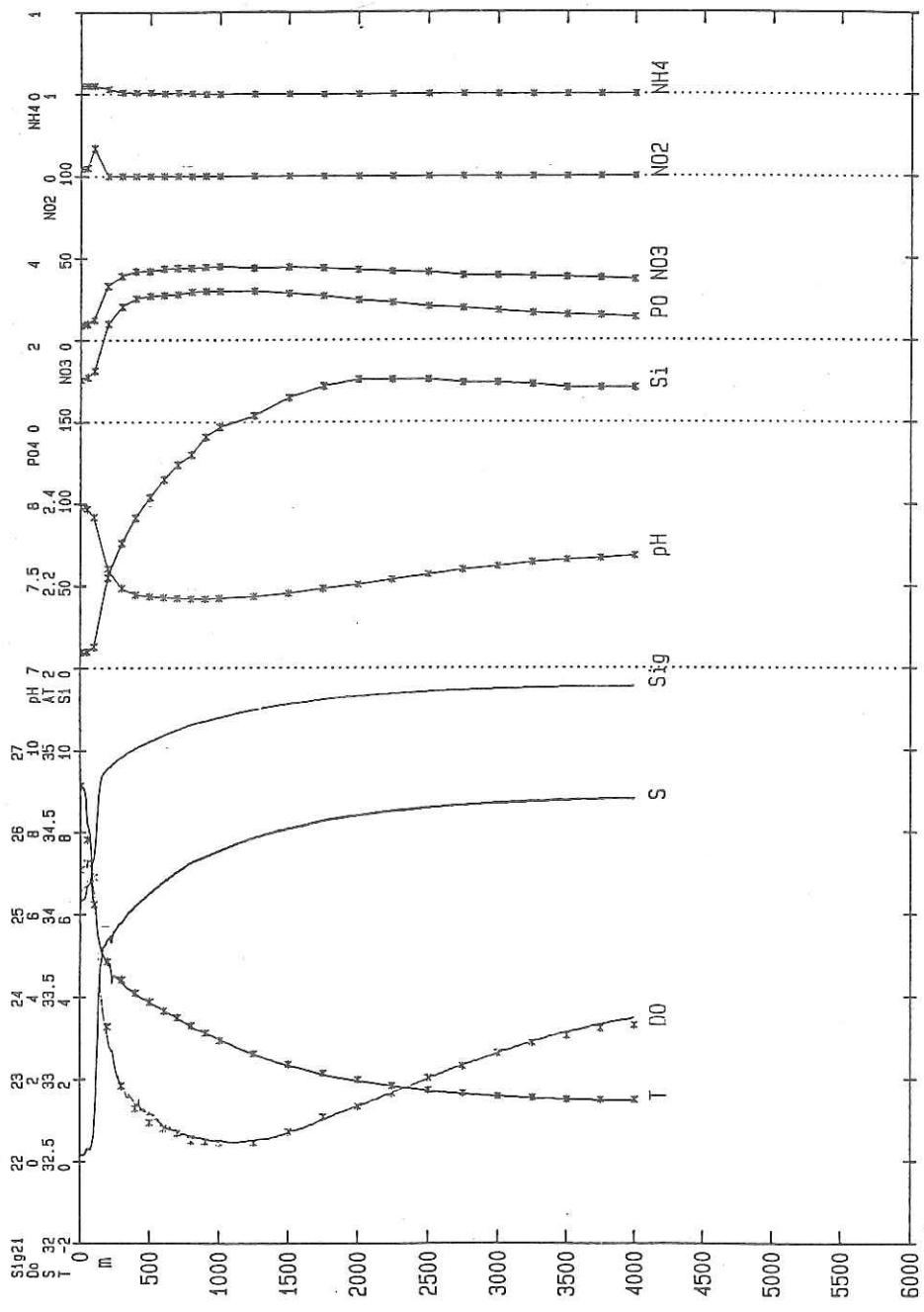


Fig. 7. Vertical profiles of temperature, salinity, sigma-t, dissolved oxygen, pH, alkalinity, silicic acid, phosphate, nitrate, nitrite and ammonium at Station G.

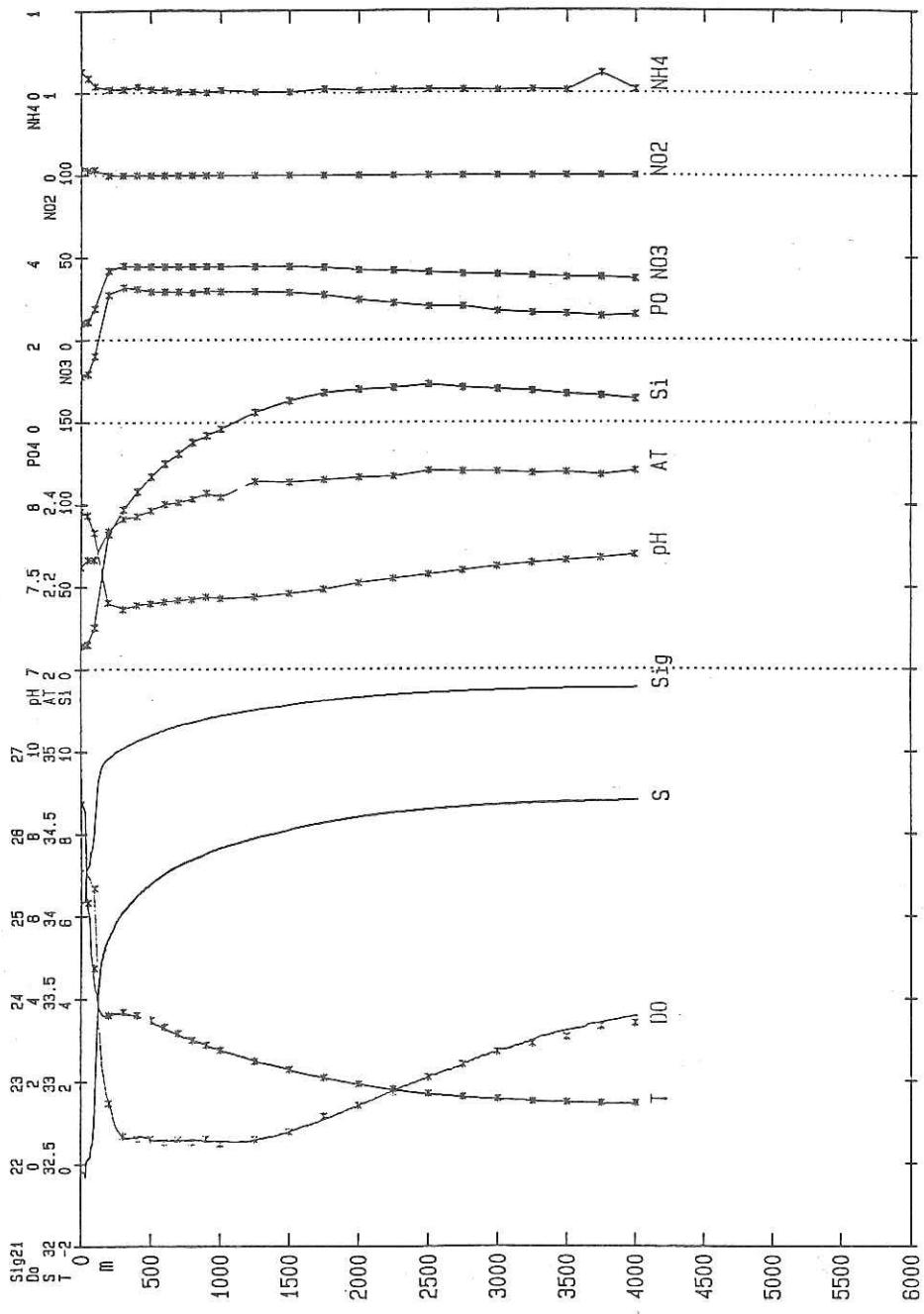


Fig. 8. Vertical profiles of temperature, salinity, sigma-t, dissolved oxygen, pH, alkalinity, silicic acid, phosphate, nitrate, nitrite and ammonium at Station F'.

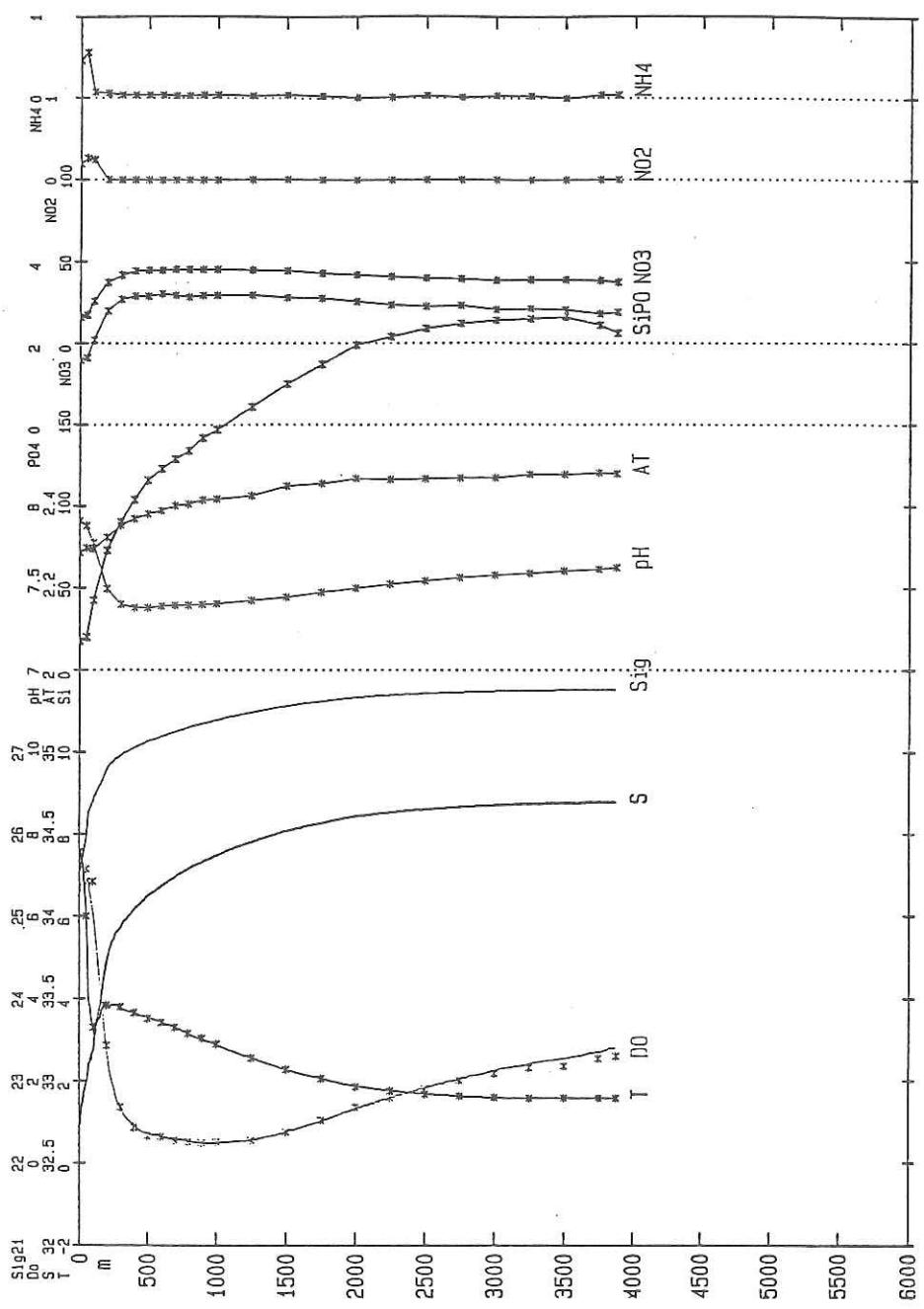


Fig. 9. Vertical profiles of temperature, salinity, sigma-t, dissolved oxygen, pH, alkalinity, silicic acid, phosphate, nitrate, nitrite and ammonium at Station C'.

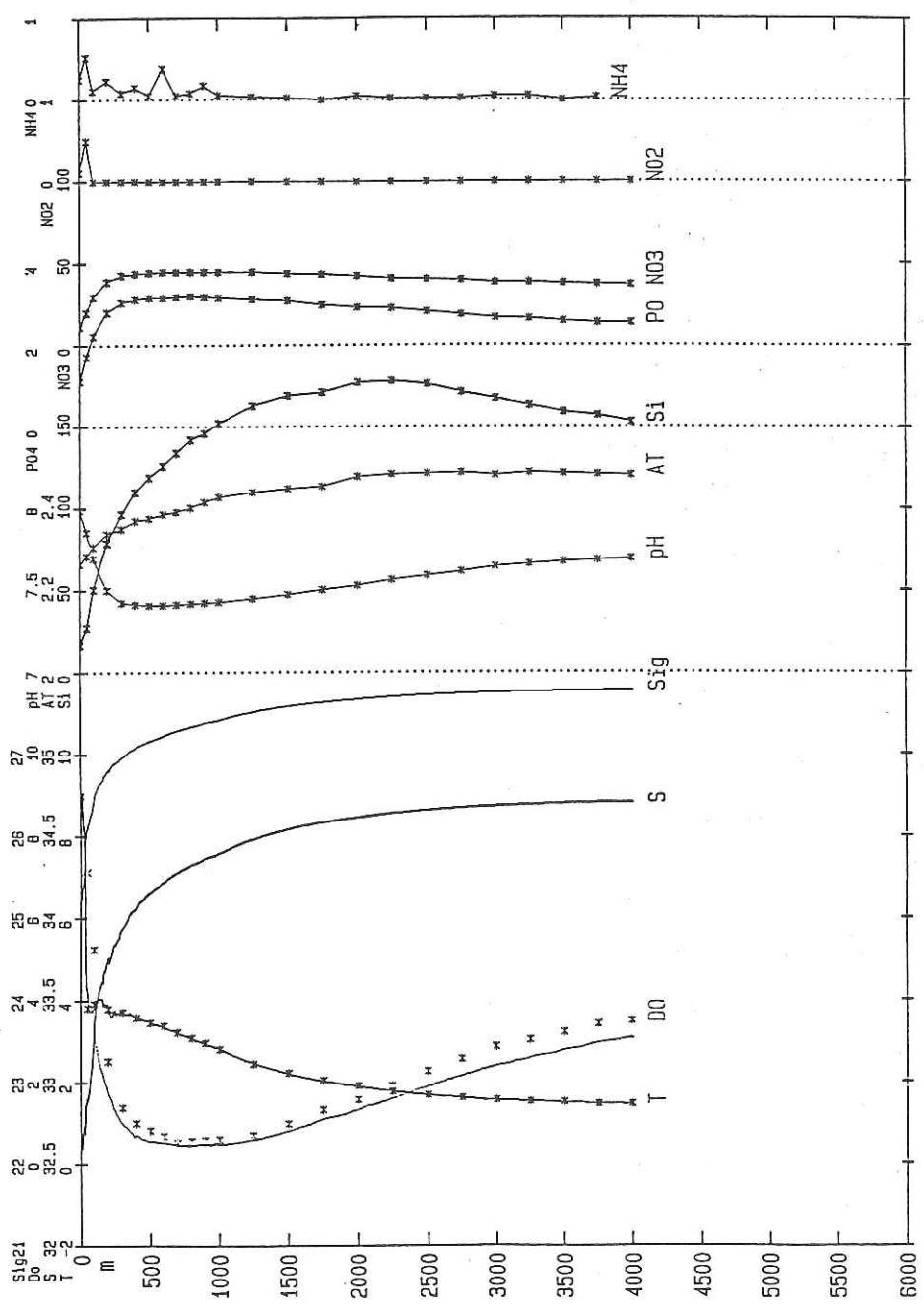


Fig. 10. Vertical profiles of temperature, salinity, sigma-t, dissolved oxygen, pH, alkalinity, silicic acid, phosphate, nitrate, nitrite and ammonium at Station H.

Table 13. Summary of hydrographic data in shallow depths at Station B

Station	B-NIS	Depth	5630 m	Date	June 9, 1986	TIME	05 : 53 - 06 : 45	Lat.	46 - 40.4 N	Long.	162 - 01.6 E	
N/S/KIN	Sample No.	D m	T C	S mg/l	D.O. mg/l	pH	AT mJ/m	S102 um	P04 um	N03 um	N02 um	NH4 um
0	0	4.20	32.996	7.81	7.867	2.355	35.6	1.92	22.1	0.29	0.26	4.200
15	15	3.84	32.999	7.77	7.869	2.326	35.2	1.99	22.1	0.29	0.32	3.839
20	20	3.84	33.004	7.83	7.869	2.302	35.8	1.87	22.4	0.29	0.43	3.839
25	24	3.79	33.007	7.74	7.866	2.330	35.6	1.91	22.4	0.29	0.46	3.786
30	29	3.58	33.013	7.71	7.860	2.312	40.3	1.92	22.6	0.28	0.47	3.576
40	39	3.42	33.052	7.62	7.863	2.303	40.9	1.98	23.4	0.28	0.44	3.410
50	49	3.28	33.067	7.78	7.821	2.302	42.0	1.99	24.5	0.29	0.48	3.277
60	59	1.74	33.417	7.72	7.808	2.299	43.2	2.03	25.1	0.32	0.45	1.737
80	78	1.47	33.133	7.72	7.799	2.302	43.6	2.09	25.6	0.40	0.44	1.466
100	98	1.49	33.191	7.19	7.761	2.321	48.3	2.22	27.2	0.47	0.38	1.425
150	146	2.86	33.656	2.61	7.465	2.342	80.7	2.92	40.2	0.06	0.09	2.851
200	3.34	33.858	1.11	7.375	2.407	94.3	3.23	44.8	0.06	0.08	3.327	26.94

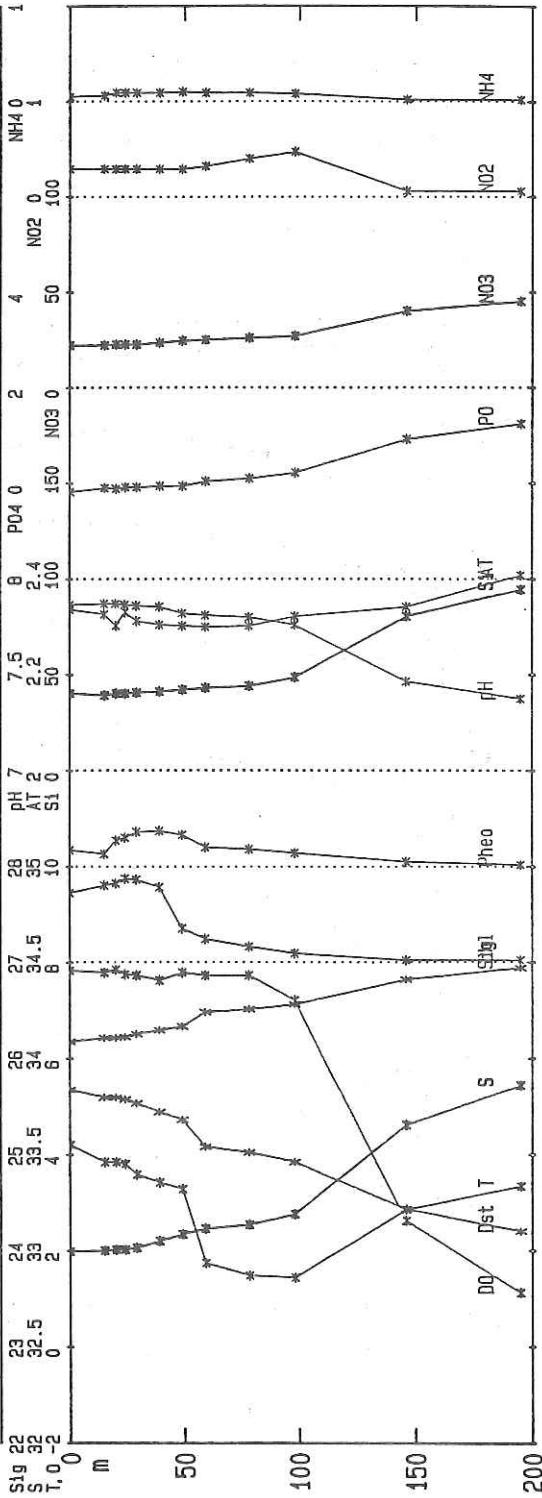


Table 14. Summary of hydrographic data in shallow depths at Station C

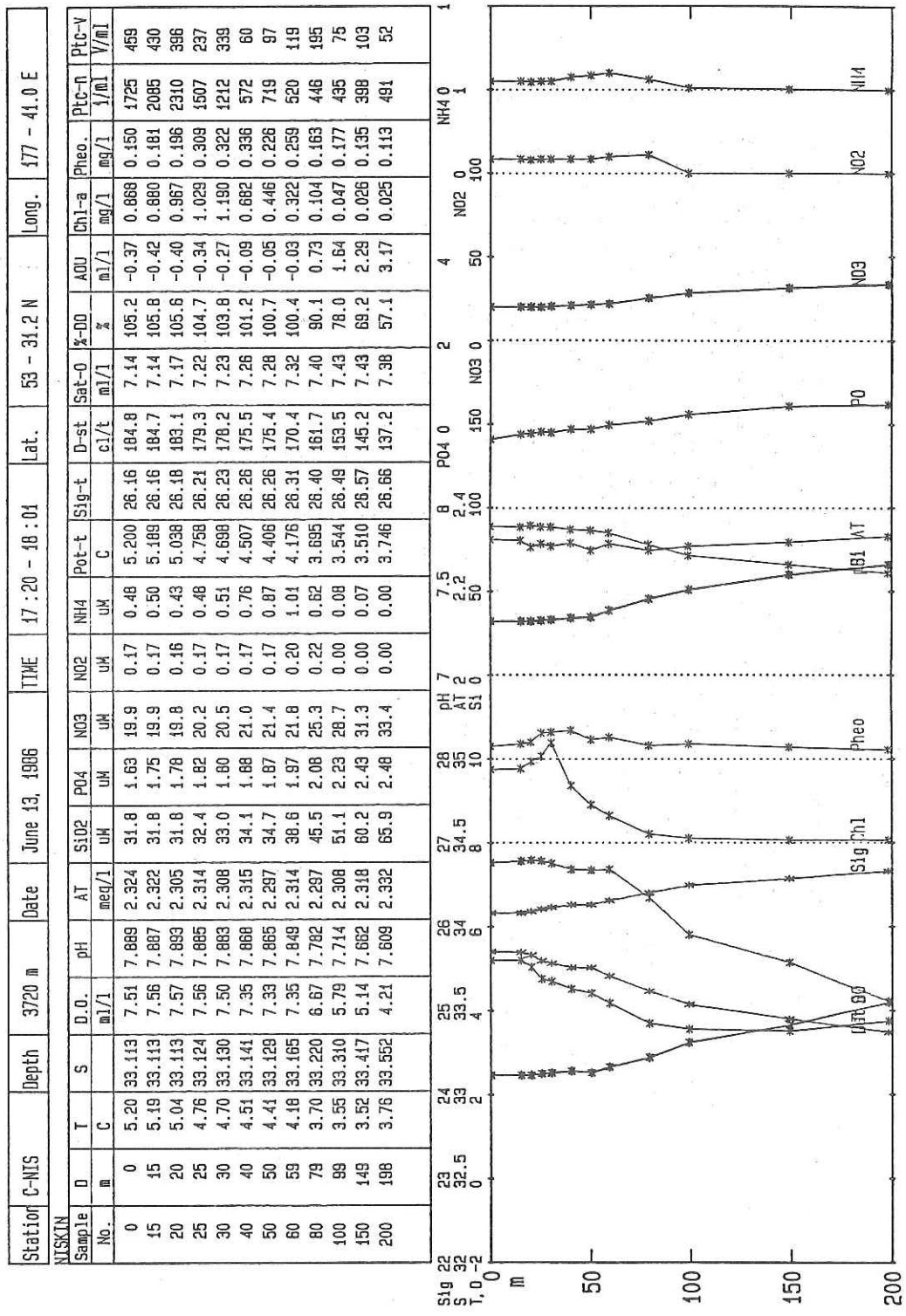


Table 15. Summary of hydrographic data in shallow depths at Station E

Station	E-NIS	Depth	3910 m	Date	June 20, 1986	TIME	09 : 01 - 09 : 48	Lat.	56 - 00.9 N	Long.	145 - 01.6 W											
WISKIN	Sample No.	D m	T C	S	D.O. ml/l	pH	AT deg/l	S102 um	P04 um	N03 um	NH4 um	Pot-t C	Sig-t	D-st cl/t	Sat-d m/l	%-d %	AOU ml/l	Chl-a mg/l	Pheo mg/l	Ptc-n 1/m	Ptc-v 1/m	
0	0	0	8.40	32.600	7.08	7.961	---	15.6	1.49	11.7	0.42	0.44	0.400	25.33	263.3	6.65	106.5	-0.43	0.645	0.097	1027	353
10	10	8.27	32.603	7.16	7.963	---	15.6	1.49	11.7	0.42	0.42	0.42	0.269	25.35	261.2	6.67	107.4	-0.49	0.422	0.102	1121	702
20	19	8.15	32.599	7.19	7.961	---	15.0	1.21	11.5	0.42	0.10	0.148	25.37	259.8	6.69	107.5	-0.50	0.558	0.126	1128	731	
25	24	7.46	32.607	7.44	7.966	---	15.0	1.19	11.7	0.42	0.17	0.458	25.47	249.9	6.79	109.5	-0.65	0.645	0.170	1181	451	
30	29	7.16	32.616	7.44	7.959	---	15.6	1.22	11.9	0.42	0.10	7.157	25.52	245.3	6.84	108.8	-0.60	0.732	0.243	1355	621	
40	39	6.60	32.624	7.36	7.943	---	16.7	1.34	12.6	0.42	0.07	6.587	25.60	237.9	6.93	106.5	-0.45	0.508	0.233	783	298	
50	49	6.26	32.615	7.24	7.928	---	17.9	1.31	13.8	0.42	0.26	0.256	25.64	234.2	6.98	103.6	-0.25	0.409	0.304	650	748	
60	58	5.65	32.632	7.25	7.914	---	18.4	1.37	14.3	0.44	0.38	5.645	25.72	225.9	7.09	102.3	-0.16	0.310	0.213	539	188	
80	78	4.96	32.754	7.06	7.882	---	20.1	1.51	15.7	0.25	0.54	4.954	25.90	209.2	7.20	98.1	0.14	0.179	0.111	420	541	
100	97	4.44	33.259	5.92	7.791	---	31.2	1.77	22.7	0.05	0.00	4.433	26.36	165.9	7.27	81.5	1.35	0.078	0.057	350	105	
150	146	3.82	33.816	1.62	7.429	---	77.0	3.00	42.6	0.00	0.00	3.810	26.86	117.6	7.35	22.0	5.73	0.018	0.025	195	77	
200	194	3.81	33.916	0.75	7.323	---	88.2	3.24	45.8	0.00	0.00	3.797	26.94	110.2	7.35	10.2	6.60	0.017	0.027	124	118	

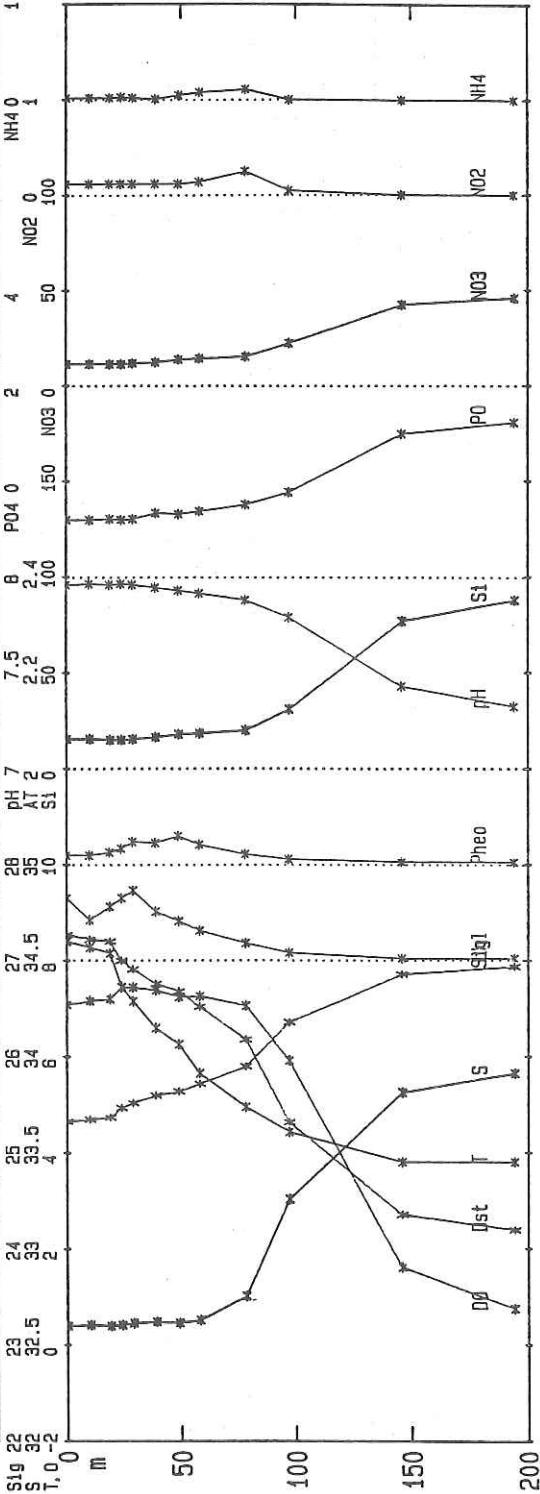


Table 16. Summary of hydrographic data in shallow depths at Station F

Station F-NIS	Depth	4100 m	Date	June 29, 1986	TIME	14:07 - 14:37	Lat.	53 - 02.9 N	Long.	144 - 59.6 W	
Sample No.	D m	T C	S mg/l	DO mg/l	pH	AT deg C	S102 um	P04 um	N03 um	N02 um	NH4 um
0	0	8.40	32.454	7.19	7.949	2.241	14.9	1.14	10.7	0.15	0.12
10	10	8.06	32.457	7.17	7.953	2.240	14.9	1.16	11.2	0.15	0.14
20	20	7.34	32.504	7.32	7.951	2.236	15.5	1.16	11.2	0.15	0.13
25	24	6.35	32.520	7.39	7.949	2.236	15.5	1.18	11.2	0.15	0.12
30	29	6.74	32.522	7.40	7.949	2.240	15.5	1.17	11.3	0.15	0.12
40	39	6.52	32.526	7.42	7.942	2.251	15.5	1.18	11.5	0.15	0.16
50	49	6.35	32.524	7.39	7.939	2.252	15.5	1.23	11.6	0.15	0.16
60	59	6.24	32.531	7.35	7.936	2.255	16.0	1.22	12.1	0.16	0.19
80	78	5.74	32.548	7.34	7.922	2.243	16.6	1.28	12.8	0.18	0.50
100	98	4.64	32.636	7.18	7.879	2.267	20.5	1.44	15.2	0.34	0.37
150	146	3.73	33.580	3.12	7.533	2.310	65.3	2.68	37.3	0.00	0.03
200	195	3.67	33.814	1.57	7.419	2.332	82.4	3.03	43.4	0.00	0.03

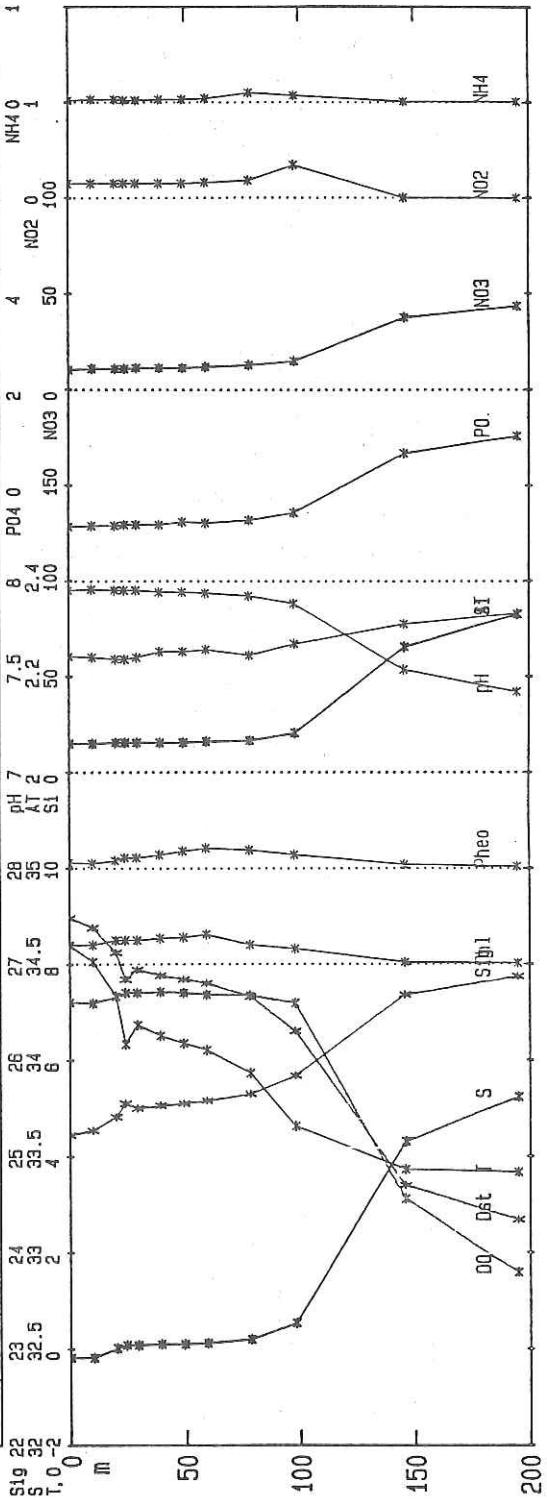


Table 17. Summary of hydrographic data in shallow depths at Station G

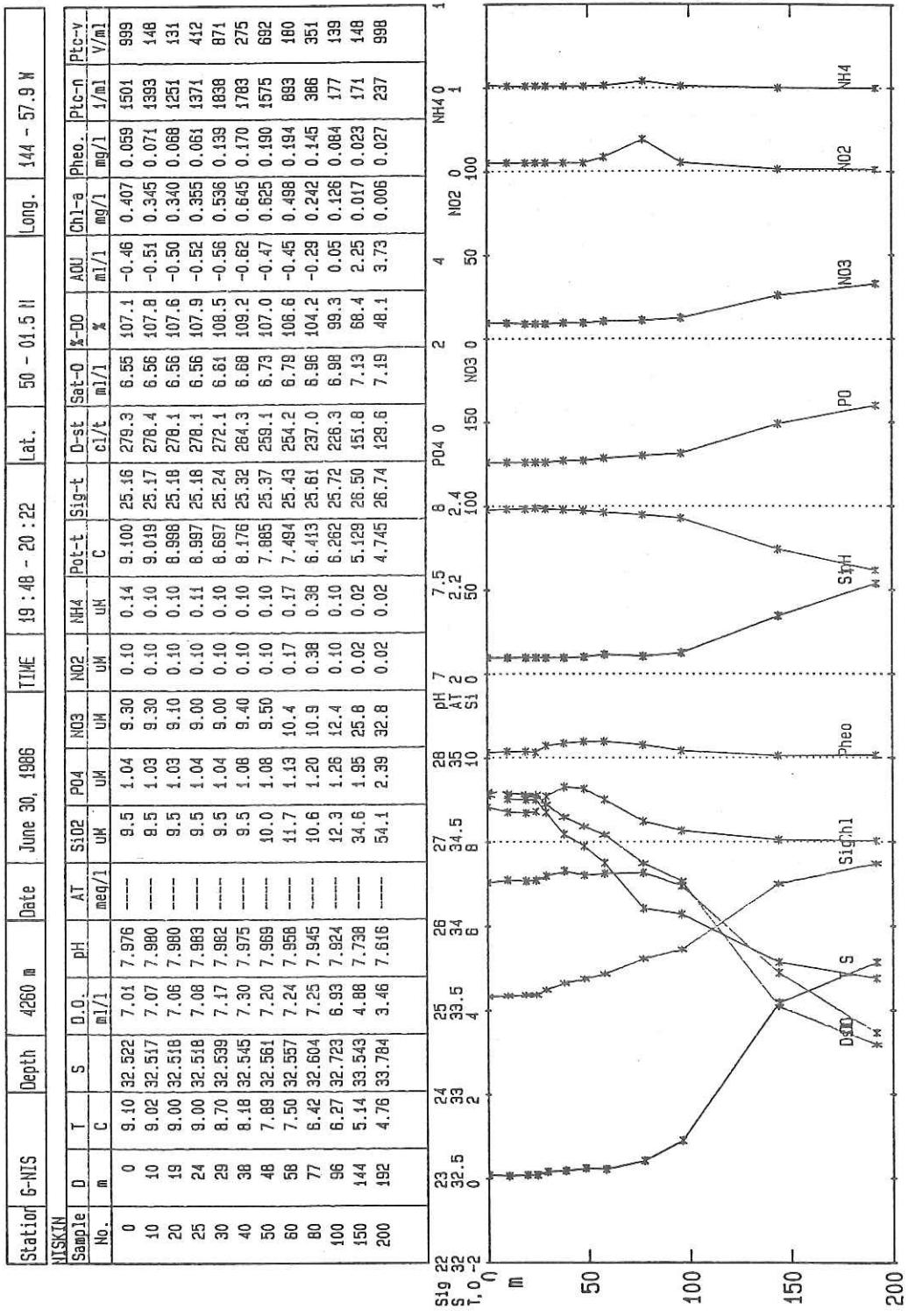


Table 18. Summary of hydrographic data in shallow depths at Station F'

Station F'-NIS	Depth	4100 m	Date	July 15, 1986	TIME	02:06 - 02:46	Lat.	53° 01.0' N	Long.	144° 56.2' W												
NISKIN	Sample	D	T	S	D.O.	pH	AT	S102	P04	N03	N02	NH4	Pot-t	Sigt	D-st	Sat-D	%-D0	AOU	Chl-a	Pheo.	Ptc-n	Ptc-y
No.	No.	m	C	m1/l	mg/l	um	um	um	um	um	um	um	C	cl/t	m1/l	%	m1/l	mg/l	mg/l	mg/l	V/m1	
0	0	8.90	32.439	7.05	7.961	2.241	13.9	4.42	10.0	0.07	0.27	8.900	25.13	282.5	6.56	107.1	-0.47	0.434	0.064	1643	197	
10	10	8.77	32.437	7.11	7.960	2.251	13.9	4.13	10.0	0.07	0.23	8.769	25.45	280.7	6.60	107.7	-0.51	0.360	0.051	1513	610	
20	20	8.71	32.446	7.12	7.956	2.251	13.9	4.15	10.2	0.08	0.23	8.708	25.16	279.2	6.61	107.7	-0.51	0.355	0.047	1204	362	
25	25	8.66	32.441	7.13	7.956	2.249	13.9	4.17	10.2	0.08	0.26	8.857	25.14	281.8	6.59	108.3	-0.54	0.353	0.040	1055	220	
30	30	8.64	32.443	7.10	7.956	2.249	13.9	4.17	10.0	0.07	0.24	8.637	25.17	278.4	6.62	107.3	-0.46	0.355	0.070	1309	691	
40	39	8.58	32.443	7.12	7.957	2.254	13.9	4.46	10.2	0.07	0.22	8.576	26.18	277.6	6.63	107.4	-0.49	0.409	0.114	1295	168	
50	49	6.95	32.507	7.42	7.944	2.250	15.6	1.21	10.7	0.07	0.26	6.946	25.46	250.8	6.88	107.9	-0.54	0.372	0.108	1221	385	
60	59	6.28	32.516	7.48	7.930	2.253	15.0	1.42	11.4	0.09	0.18	6.275	25.56	241.9	6.99	107.1	-0.49	0.360	0.164	772	558	
80	79	5.41	32.556	7.37	7.910	2.258	16.2	1.34	12.7	0.18	0.50	5.404	25.59	226.8	7.43	103.3	-0.24	0.302	0.200	428	124	
100	99	4.43	32.763	6.65	7.830	2.270	24.9	1.61	18.6	0.06	0.10	4.423	25.96	203.2	7.29	91.2	0.64	0.151	0.121	277	44	
150	148	3.71	33.641	2.65	7.490	2.327	68.0	2.83	37.6	0.00	0.10	3.700	26.73	130.0	7.38	35.9	4.73	0.047	0.037	301	22	
200	197	3.60	33.825	1.54	7.411	2.346	81.6	3.08	42.6	0.00	0.07	3.567	26.89	115.1	7.39	20.8	5.85	0.010	0.026	217	128	

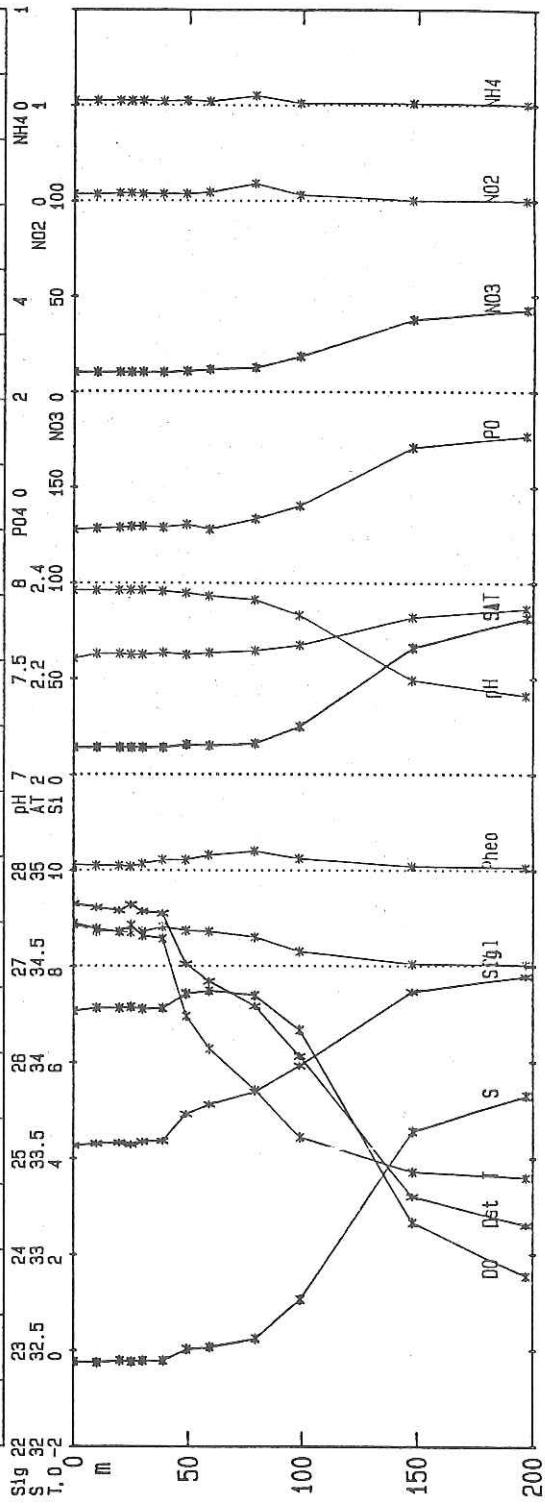


Table 19. Summary of hydrographic data in shallow depths at Station C'

Station	C'-NIS	Depth	3930 m	Date	July 22, 1986	TIME	15:16 - 16:00	Lat.	53° - 30.5 N	Long.	177° - 32.1 E
NISKIN	Sample	O	T	S	D.O.	pH	AT	SI:02	PO4	NO3	NH4
No.	m	C		ml/l	mg/l	meq/l	°C	UW	UW	UW	UW
0	0	7.55	32.906	7.14	7.904	2.286	16.3	1.12	45.7	0.48	0.47
10	10	7.42	32.910	7.14	7.906	2.290	16.3	1.13	45.7	0.48	0.44
20	20	7.26	32.917	7.14	7.906	2.274	16.3	1.51	45.8	0.48	0.49
25	24	7.15	32.930	7.18	7.908	2.282	16.3	1.51	45.8	0.48	0.56
30	29	7.04	32.951	7.17	7.909	2.294	17.6	1.49	46.0	0.17	0.67
40	39	6.55	32.976	7.13	7.899	2.296	18.8	1.58	46.6	0.22	0.56
50	49	5.93	33.004	7.45	7.874	2.290	20.1	1.65	18.1	0.27	0.64
60	59	5.32	33.049	7.39	7.868	2.303	23.9	1.76	19.1	0.28	0.76
80	78	3.75	33.125	7.09	7.793	2.284	33.9	2.00	23.1	0.27	0.40
100	98	3.32	33.179	6.78	7.759	2.287	40.2	2.14	26.8	0.03	0.04
150	146	3.50	33.299	6.70	7.724	2.302	52.8	2.29	29.8	0.00	0.07
200	195	3.86	33.664	2.83	7.492	2.318	71.6	2.76	38.4	0.00	0.09

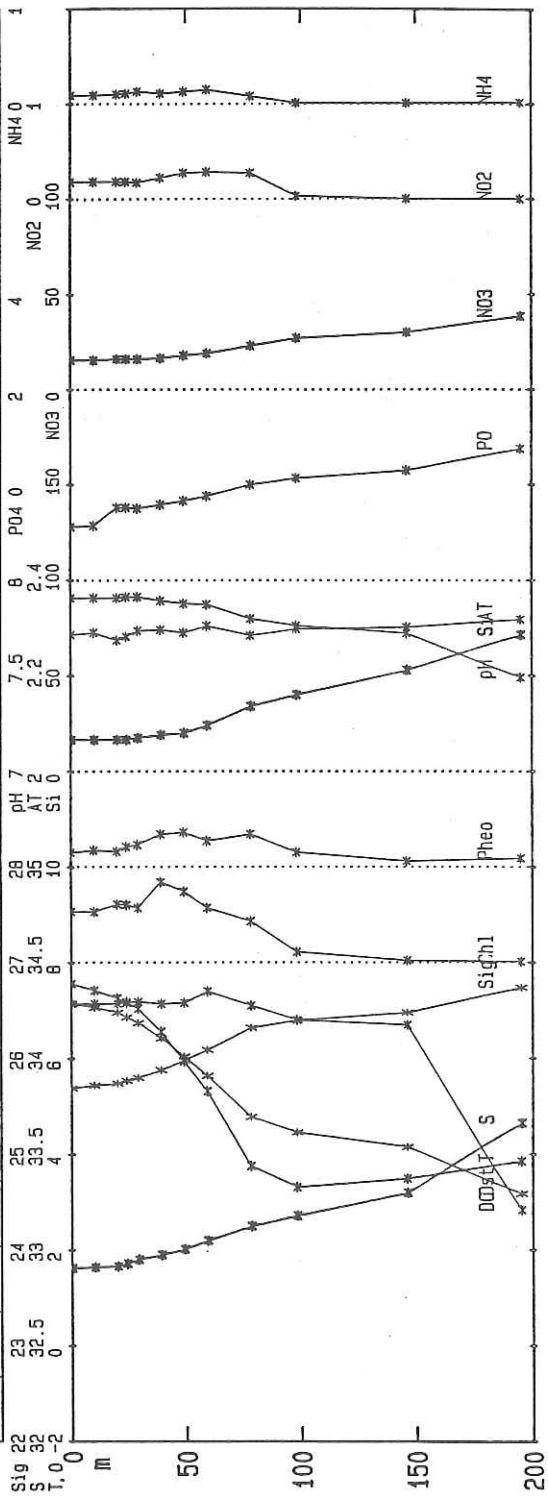


Table 20. Summary of hydrographic data in shallow depths at Station H

Station	H-NIS	Depth	4820 m	Date	July 25, 1986	TIME	03:37 - 04:13	Lat.	51° 05.0' N	Long.	170° 56.8' E									
NISKIN Sample No.	D C	T	S	0.0. mg/l	pH	AT	S102 UM	P04 UM	N02 UM	NH4 UM	Po-t C	Sig-t	D-st cl/t	Sat-d m1/l	%-d %	AOU m1/l	Chl-a mg/l	Pheo. mg/l	Ptc-y V/ml	
0	0	9.00	32.595	7.01	7.991	2.263	16.5	1.42	11.1	0.11	0.28	9.000	25.24	272.3	6.56	106.9	-0.45	1.017	0.176	3686 1472
10	10	8.25	32.689	7.14	7.989	2.272	17.7	1.43	11.6	0.11	0.26	8.249	25.42	254.5	6.67	107.1	-0.47	1.012	0.304	3972 1502
20	19	7.69	32.757	7.17	7.976	2.275	18.9	1.25	12.3	0.12	0.38	7.688	25.56	241.8	6.75	106.2	-0.42	0.990	0.533	3088 556
25	24	6.45	32.860	7.32	7.938	2.274	22.4	1.42	14.8	0.17	0.48	6.448	25.81	218.2	6.94	105.4	-0.38	1.078	0.522	2821 562
30	29	5.58	32.893	7.26	7.905	2.279	23.6	1.52	16.4	0.24	1.00	5.578	25.94	205.5	7.09	102.4	-0.17	0.990	0.558	1971 523
40	38	4.50	32.876	7.25	7.881	2.282	23.6	1.62	17.6	0.45	1.04	4.487	26.05	195.3	7.28	99.7	0.03	0.521	0.305	836 363
50	48	4.09	32.886	7.18	7.866	2.284	24.8	1.70	18.9	0.59	1.40	4.087	26.10	190.6	7.35	97.7	0.47	0.270	0.174	741 4356
60	58	3.81	32.925	7.01	7.840	2.285	28.5	1.76	21.4	0.29	0.12	3.806	26.15	185.0	7.40	94.8	0.39	0.139	0.145	614 62
80	79	3.78	33.094	6.41	7.767	2.283	38.9	2.01	25.5	0.00	0.14	3.775	26.29	172.0	7.39	82.6	1.28	0.056	0.091	522 369
100	96	3.84	33.268	5.26	7.700	2.308	48.3	2.18	29.3	0.00	0.14	3.894	26.42	159.4	7.38	71.3	2.42	0.052	0.076	409 141
150	144	4.08	33.535	3.95	7.605	2.327	66.0	2.50	34.0	0.00	0.12	4.070	26.61	141.5	7.32	54.0	3.37	0.011	0.080	494 50
200	192	3.82	33.714	2.63	7.506	2.332	75.5	2.62	38.6	0.00	0.17	3.807	26.78	125.5	7.36	35.7	4.73	0.005	0.069	856 131
Sig 22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	
Sig 32	32.5	33.5	33.5	34	34.5	34.5	35	35	35	35	35	35	35	35	35	35	35	35	35	
T	-2	0	2	4	6	8	10	12	14	16	18	20	22	24	26	28	30	32	34	
m	0	50	100	150	200	250	300	350	400	450	500	550	600	650	700	750	800	850	900	

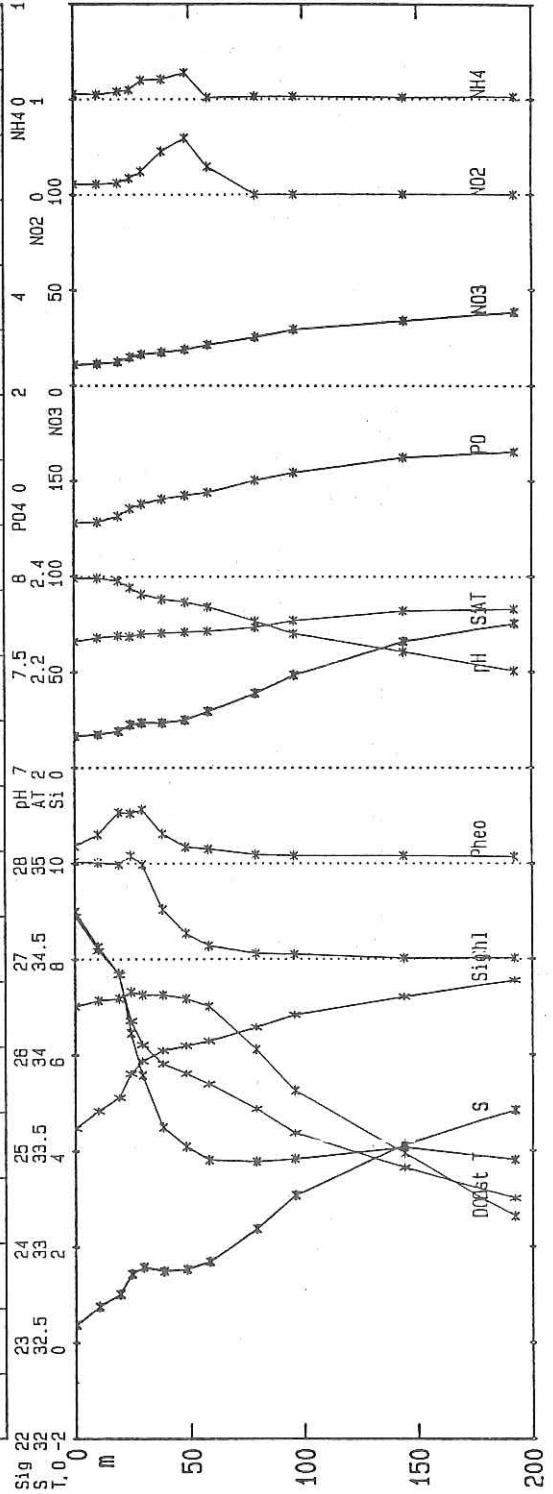


Table 21. Summary of OCTOPUS data at Station C

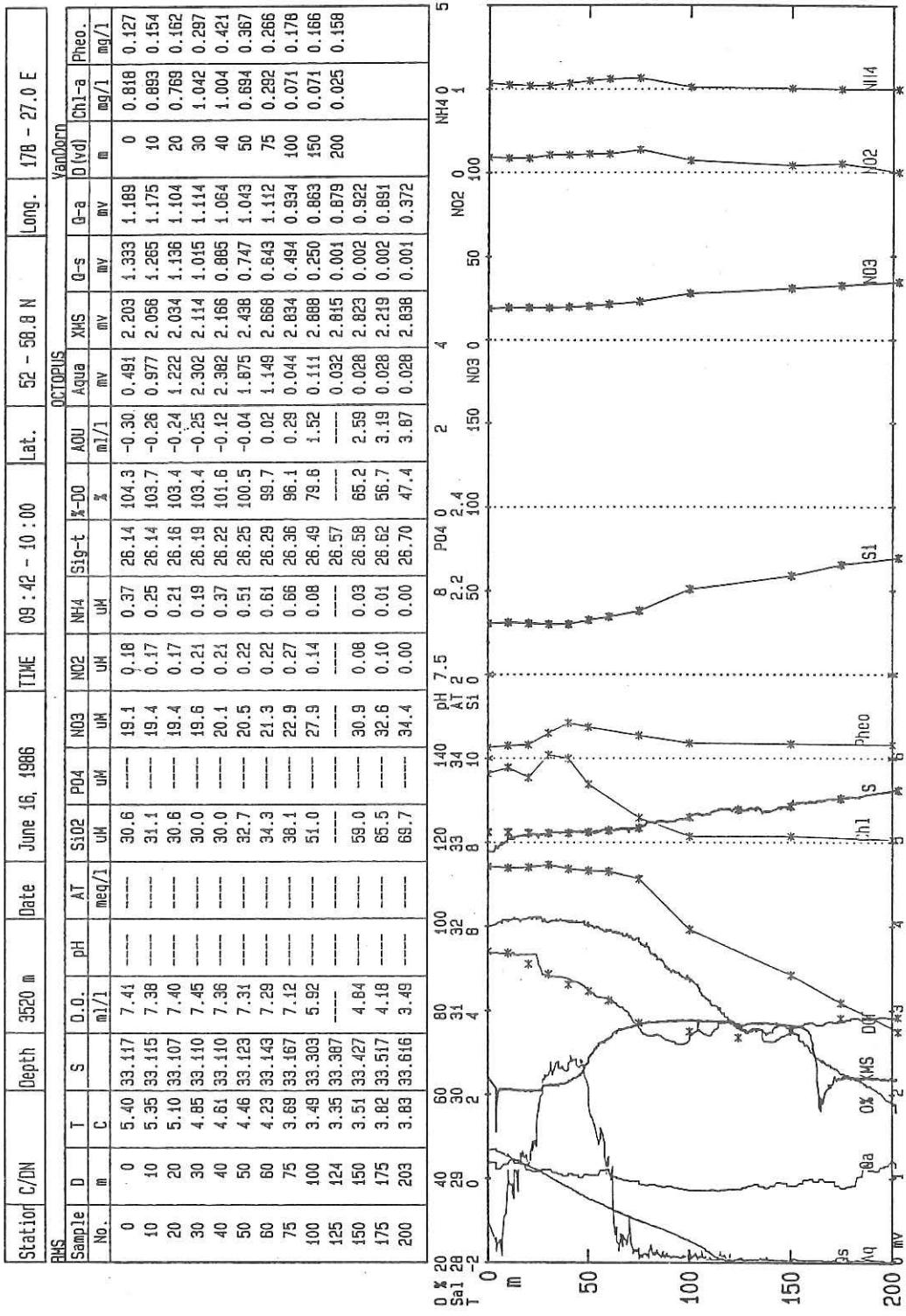


Table 22. Summary of OCTOPUS data at Station 1

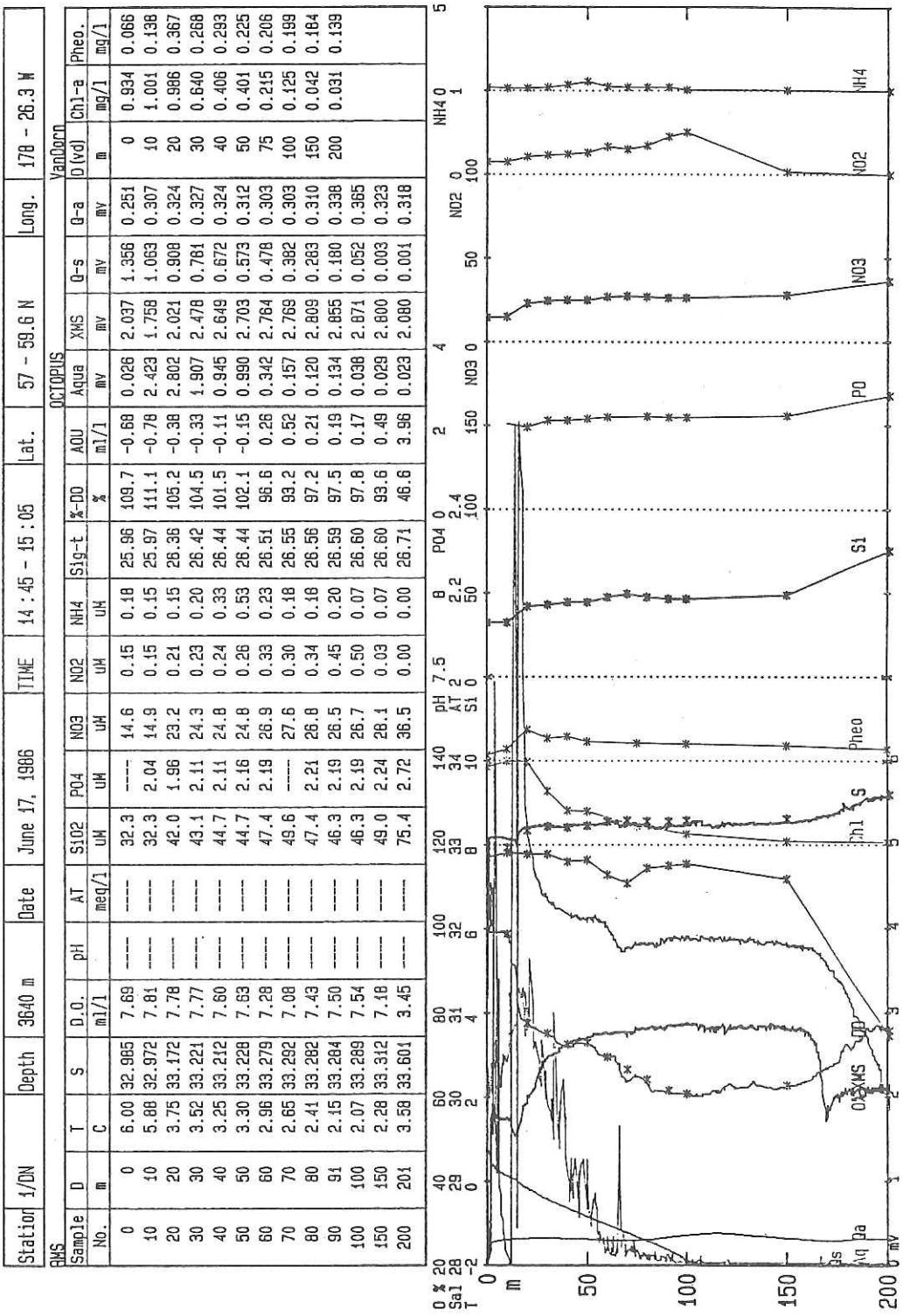


Table 23. Summary of OCTOPUS data at Station 2

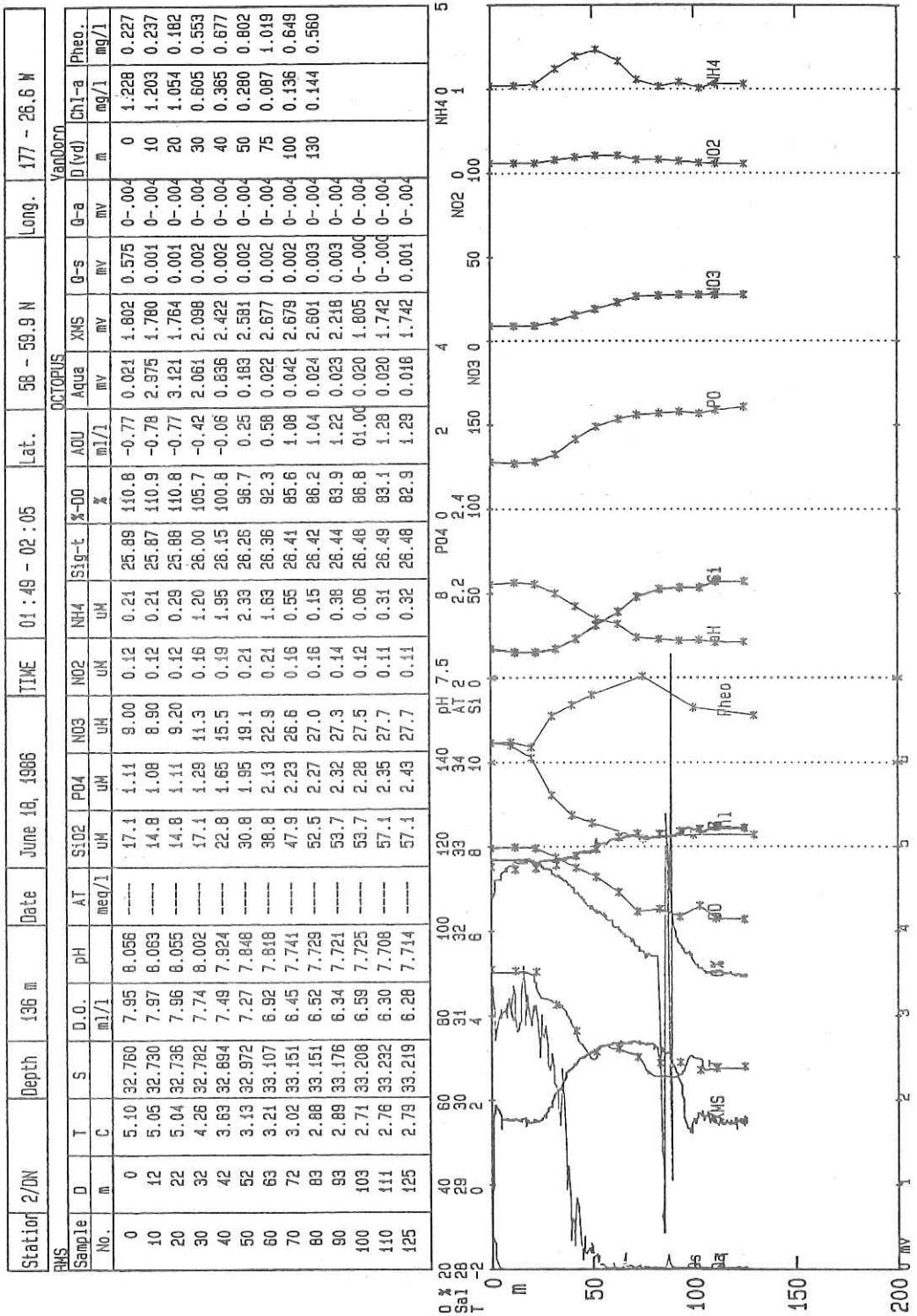


Table 24. Summary of OCTOPUS data at Station 3

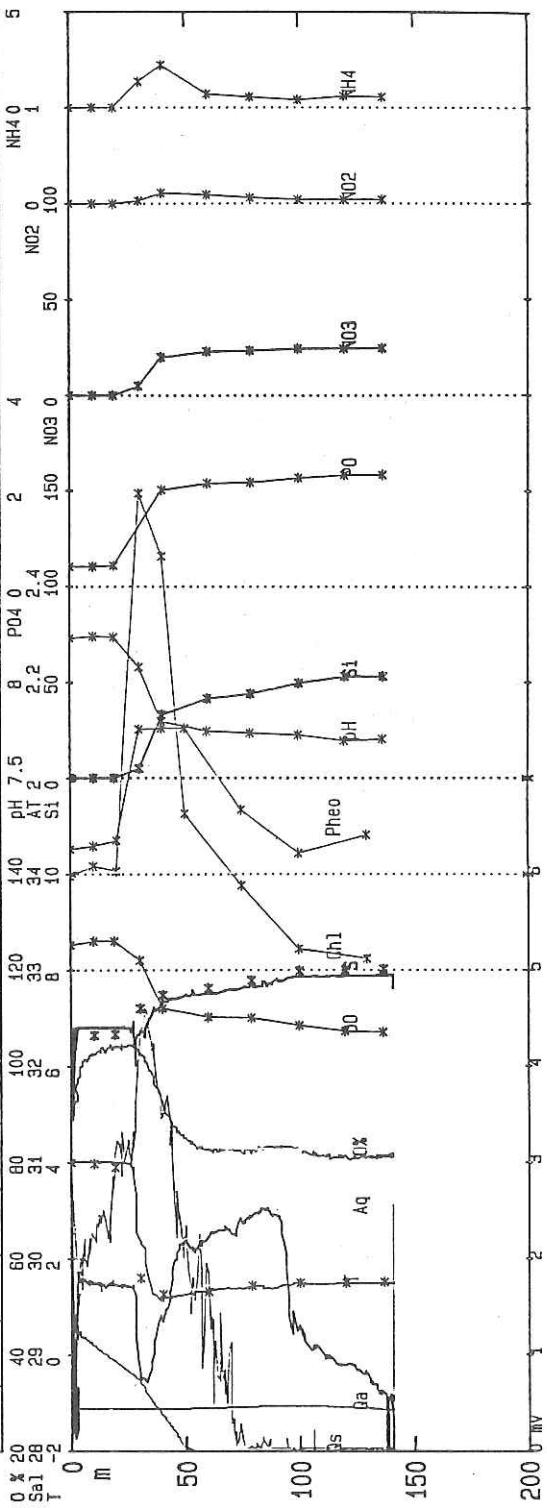


Table 25. Summary of OCTOPUS data at Station 4

Station	4/DN	Depth	140 m	Date	June 18, 1986	TIME	16:50 - 17:08	Lat.	60° - 40.7 N	Long.	175° - 30.2 W										
AMS Sample No.	D C	T	S	D.O.	pH	AT	S102	PO4	N03	N02	NH4	Sig-t %	ADU ml/l	Aqua mv	XMS mv	G-s mv	G-a mv	Vandboro D(vd) mg/l	Chl-a Pheo. mg/l		
0	0	3.20	31.200	8.05	7.990	---	0.0	0.39	0.0	0.00	0.01	24.84	106.0	-0.45	1.534	2.338	1.494	0.720	0	0.229	0.060
.5	6	3.24	31.216	8.12	7.995	---	0.0	0.41	0.0	0.00	0.00	24.85	107.0	-0.53	0.044	2.424	1.275	0.703	10	0.234	0.058
15	15	2.05	31.482	8.22	8.009	---	0.0	0.39	0.0	0.00	0.00	25.45	105.3	-0.41	0.044	2.281	1.171	0.706	20	0.247	0.068
25	25	1.68	31.448	8.32	8.016	---	0.0	0.39	0.0	0.00	0.00	25.43	106.1	-0.48	0.098	2.302	1.077	0.738	30	0.368	0.096
35	35	-0.65	31.682	9.76	8.051	---	0.0	0.49	0.0	0.00	0.00	25.45	116.6	-1.39	1.482	1.547	0.973	0.806	40	1.203	0.310
45	45	-4.49	31.873	9.23	---	2.3	---	2.0	0.05	0.08	25.63	107.9	-0.68	2.350	1.818	0.793	0.878	50	0.521	0.221	
55	55	-1.36	32.046	8.44	7.877	---	17.0	1.19	9.00	0.13	0.82	25.77	99.1	0.07	0.409	2.633	0.626	0.795	75	0.496	0.319
65	65	-0.73	32.258	7.68	7.799	---	27.5	1.64	14.8	0.10	0.27	25.92	91.9	0.68	0.043	2.786	0.515	0.774	100	0.066	0.289
75	75	0.13	32.425	6.60	7.724	---	33.0	1.96	17.9	0.45	2.28	26.02	80.9	1.56	0.042	2.467	0.409	0.776			
85	85	0.54	32.505	6.35	7.663	---	41.9	---	20.1	0.14	2.38	26.07	78.7	1.71	0.043	2.004	0.275	0.740			
95	95	0.56	32.504	6.28	7.659	---	42.4	2.49	20.1	0.14	2.54	26.06	77.9	1.78	0.042	1.957	0.024	0.717			
105	105	0.56	32.501	6.26	7.659	---	42.4	2.15	20.0	0.14	2.50	26.06	77.7	1.80	0.072	2.670	0.071	0.358			

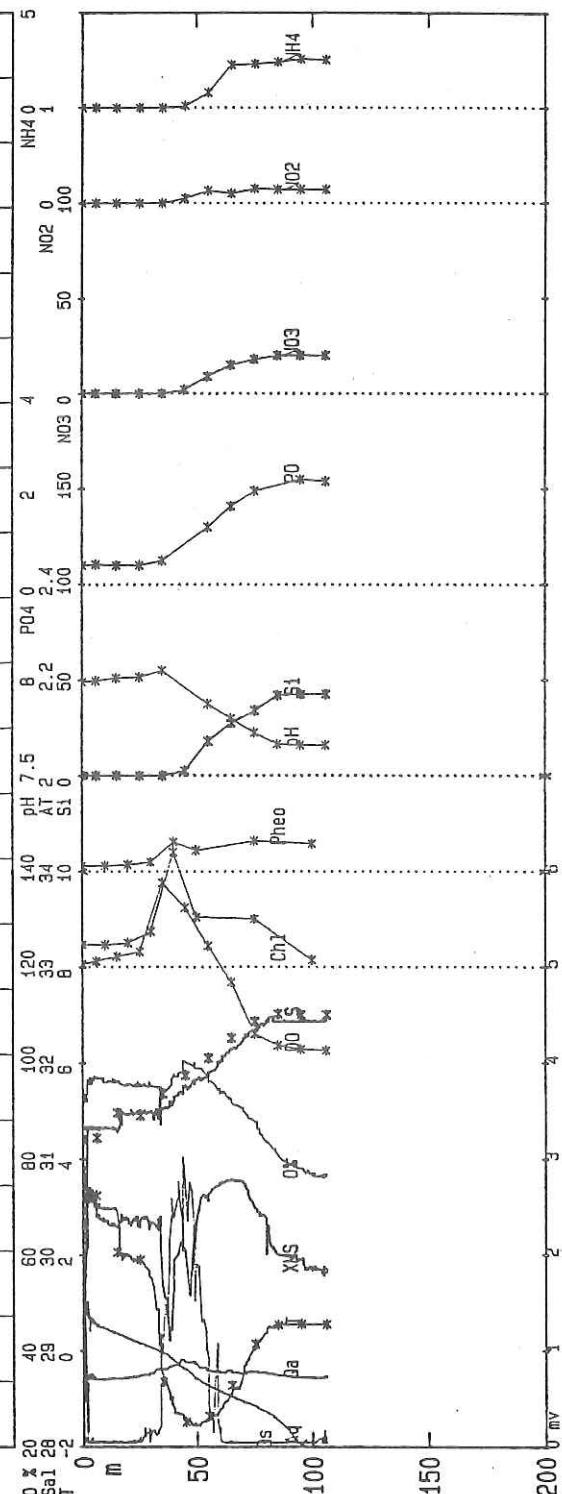


Table 26. Summary of OCTOPUS data at Station 5

Station	5/DN	Depth	79 m	Date	June 18-19, 1986	TIME	23:58 - 00:13	Lat.	61° 42.5' N	Long.	OCTOPUS																							
											D	T	S	D.O.	pH	AT	Si/02	P04	N03	N02	NH4	Sig-t	%-S0	AOU	Aqua	XMS	g-s	B-a	MV	MV	Chl-a	Phae.		
No.	m	°C		ml/l	mg/l	UN	UN	UN	UN	UN	UN	%	ml/l	mg/l	mg/l	mg/l	UN	UN	UN	UN	UN	UN	UN	mg/l	mg/l									
0	0	3.40	30.479	8.34	8.055	—	—	0.0	0.53	0.0	0.00	0.10	24.27	109.0	-0.69	0.021	2.738	0.267	0.-003	0	0.204	0.032												
5	5	3.22	30.507	8.35	8.055	—	—	0.0	0.51	0.0	0.00	0.07	24.28	109.5	0.72	0.016	2.592	0.137	0.-003	10	0.156	0.045												
10	10	1.99	30.850	9.34	9.116	—	—	0.0	0.56	0.0	0.00	0.05	24.65	148.9	-1.49	0.020	2.570	0.003	0.-003	20	0.593	0.094												
15	15	0.60	31.389	9.66	—	—	—	0.0	0.57	0.0	0.00	0.38	25.16	119.1	-1.55	0.018	2.472	0.000	0.-003	30	1.628	0.591												
20	20	0.44	31.671	9.96	8.116	—	—	0.0	0.56	0.0	0.00	0.02	25.40	122.5	-1.83	0.015	2.449	0.001	0.-003	40	1.032	0.213												
25	25	-0.99	31.865	10.52	8.078	—	—	0.5	0.63	0.0	0.00	0.00	25.69	124.7	-2.09	0.865	1.993	0.001	0.-003	50	1.086	0.256												
30	30	-1.45	32.057	10.07	7.872	—	—	1.6	0.76	0.0	0.00	0.14	25.78	118.0	-1.54	2.393	1.772	0.001	0.-003	75	1.386	0.627												
35	35	-1.62	32.156	8.64	—	—	—	4.7	1.30	0.40	0.14	0.97	25.87	100.8	-0.07	3.224	1.743	0.001	0.-003															
40	40	-1.66	32.205	8.25	7.692	—	—	12.7	1.45	9.70	0.16	1.49	25.91	96.2	0.33	0.077	2.140	0.001	0.-003															
45	45	-1.70	32.247	7.83	7.786	—	—	25.3	1.64	14.0	0.17	1.63	25.94	91.2	0.75	0.841	1.850	0.002	0.-003															
50	50	-1.70	32.266	7.83	—	—	—	24.3	1.67	11.1	0.46	1.67	25.96	91.2	0.75	0.017	1.378	0.002	0.-003															
60	60	-1.70	32.288	7.73	7.784	—	—	25.3	1.73	11.4	0.16	1.77	25.97	90.1	0.85	0.016	0.546	0.-000	0.-003															
75	75	-1.69	32.359	7.21	7.702	—	—	26.4	1.75	9.20	0.13	1.79	26.03	84.1	1.36	0.353	0.288	0.-001	0.-003															
								140	7.5	8	2	2.2	100	120	34	AT 2	2.4	0	4	NH4 0	1													
								32	31	30	29	28	20	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	
								6	5	4	3	2	1	0	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20

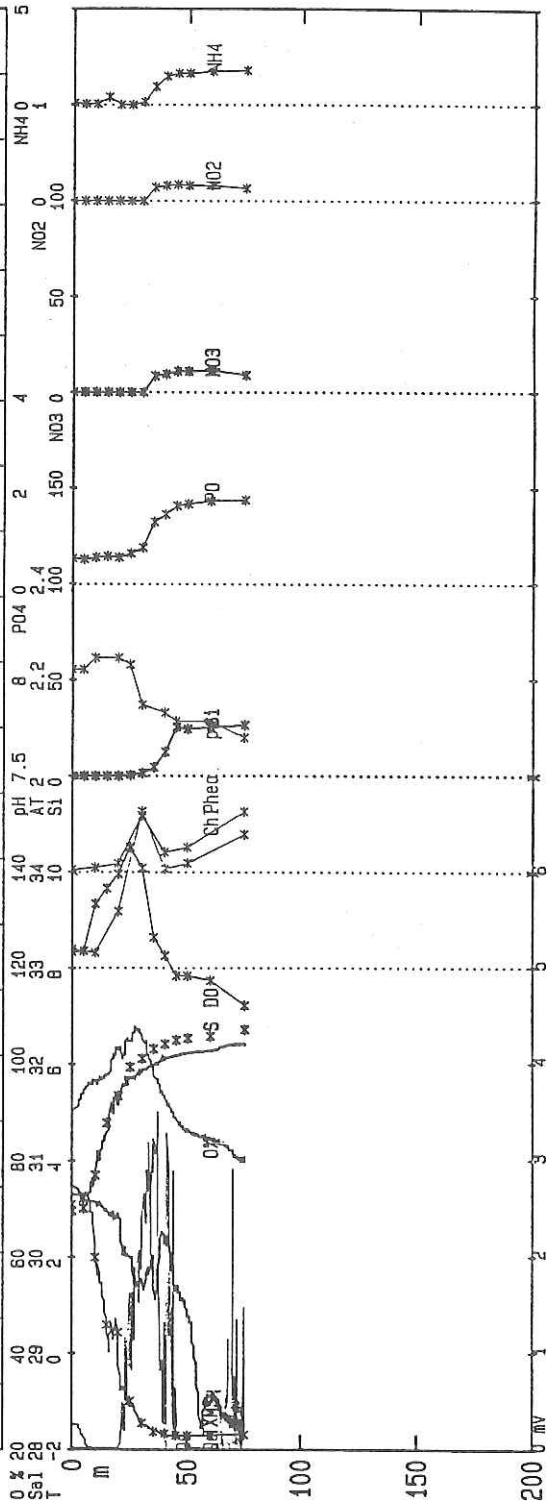


Table 27. Summary of OCTOPUS data at Station 6

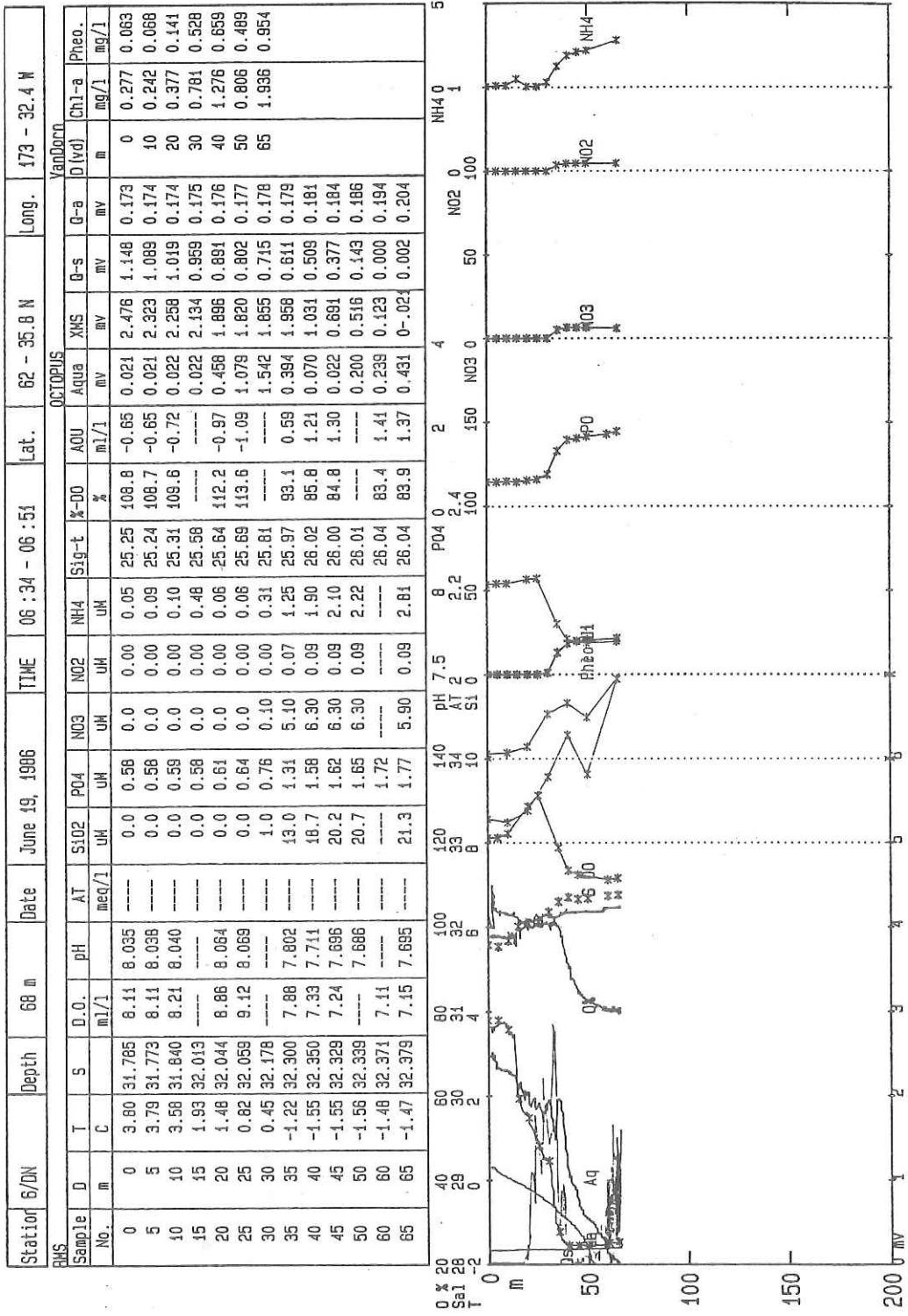


Table 28. Summary of OCTOPUS data at Station D

Station D/DN	Depth	52 m	Date	June 19, 1986	TIME	12 : 32 - 12 : 42	Lat.	63 - 30.0 N	Long.	172 - 30.0 W	OCTOPUS										
											Vandorn										
Sample No.	D	T	S	D.O.	pH	AT	SiO ₂	Po ₄	NH ₄	Sig-t %	AOU	XWS	O-s	O-a	O(kd)	Chl-a	Pheo.				
	m	C	‰	ml/l		mg/l	µM	µM	µM	%	ml/l	mv	mv	mv	mg/l	mg/l					
0	0	3.40	32.155	8.53	8.092	2.251	0.0	0.53	0.0	0.04	25.56	143.6	-1.02	0.026	2.422	1.479	0.593	0	0.244	0.113	
5	5	3.28	32.163	8.55	8.102	2.246	0.0	0.54	0.0	0.02	25.60	143.5	-1.02	0.028	2.34	1.243	0.586	10	0.290	0.112	
10	10	3.14	32.158	8.70	8.104	2.247	0.0	0.54	0.0	0.04	25.60	145.1	-1.14	0.026	2.168	1.173	0.594	20	0.588	0.265	
15	15	2.60	32.189	—	8.105	2.248	0.0	0.51	0.0	0.02	25.67	—	—	0.060	2.207	1.112	0.601	30	0.840	1.918	
20	20	2.55	32.194	9.65	8.108	2.256	0.0	0.56	0.0	0.04	25.68	125.8	-1.98	0.497	2.050	1.061	0.608	40	2.807	1.859	
25	25	-0.92	32.446	10.26	8.054	2.257	2.1	0.79	2.00	0.01	0.58	26.08	122.3	-1.87	1.471	1.909	0.989	615	50	1.144	2.108
30	30	-1.22	32.549	9.72	8.028	2.260	2.1	0.88	1.90	0.04	0.92	26.17	115.0	-1.27	3.752	1.238	0.870	621			
35	35	-1.20	32.620	—	8.027	2.269	6.7	1.01	3.40	0.07	1.95	26.23	—	—	1.563	2.328	0.705	625			
40	41	-1.35	32.806	7.24	7.717	2.282	25.9	1.92	9.00	0.13	4.34	26.39	85.5	1.23	1.437	0.695	0.624	624			
45	45	-1.35	32.806	7.10	7.707	2.280	25.9	1.93	9.30	0.13	4.35	26.39	83.9	1.37	1.614	0.319	0.499	622			

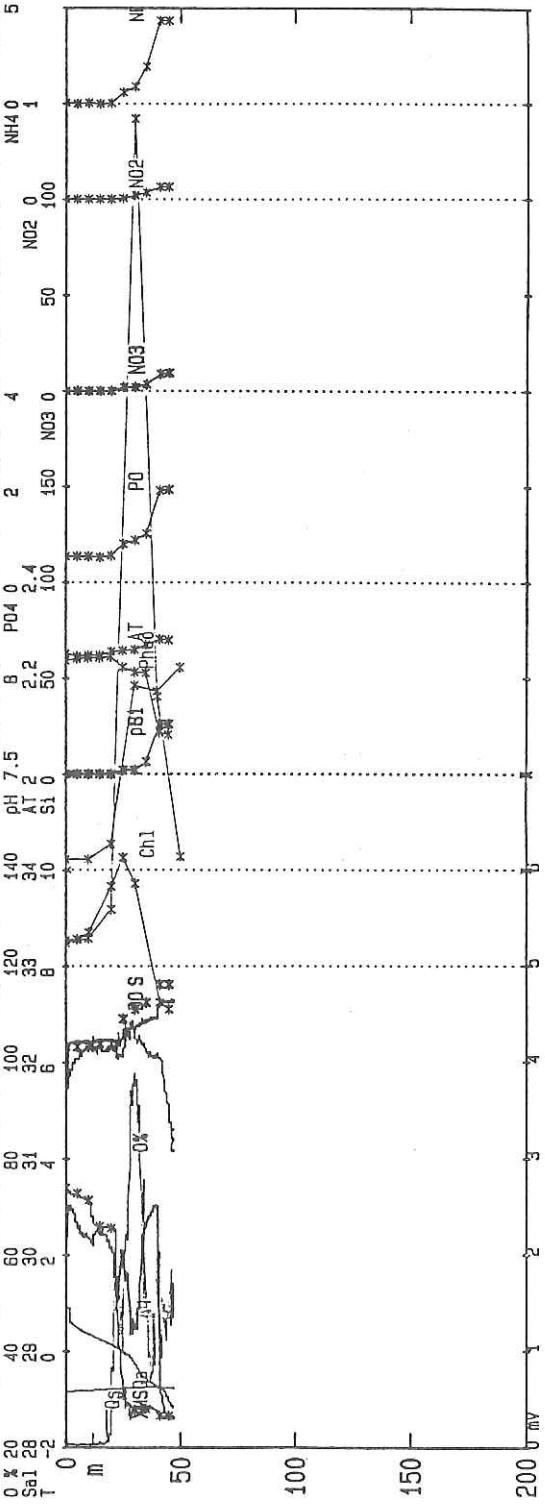


Table 29. Summary of OCTOPUS data at Station 7-1

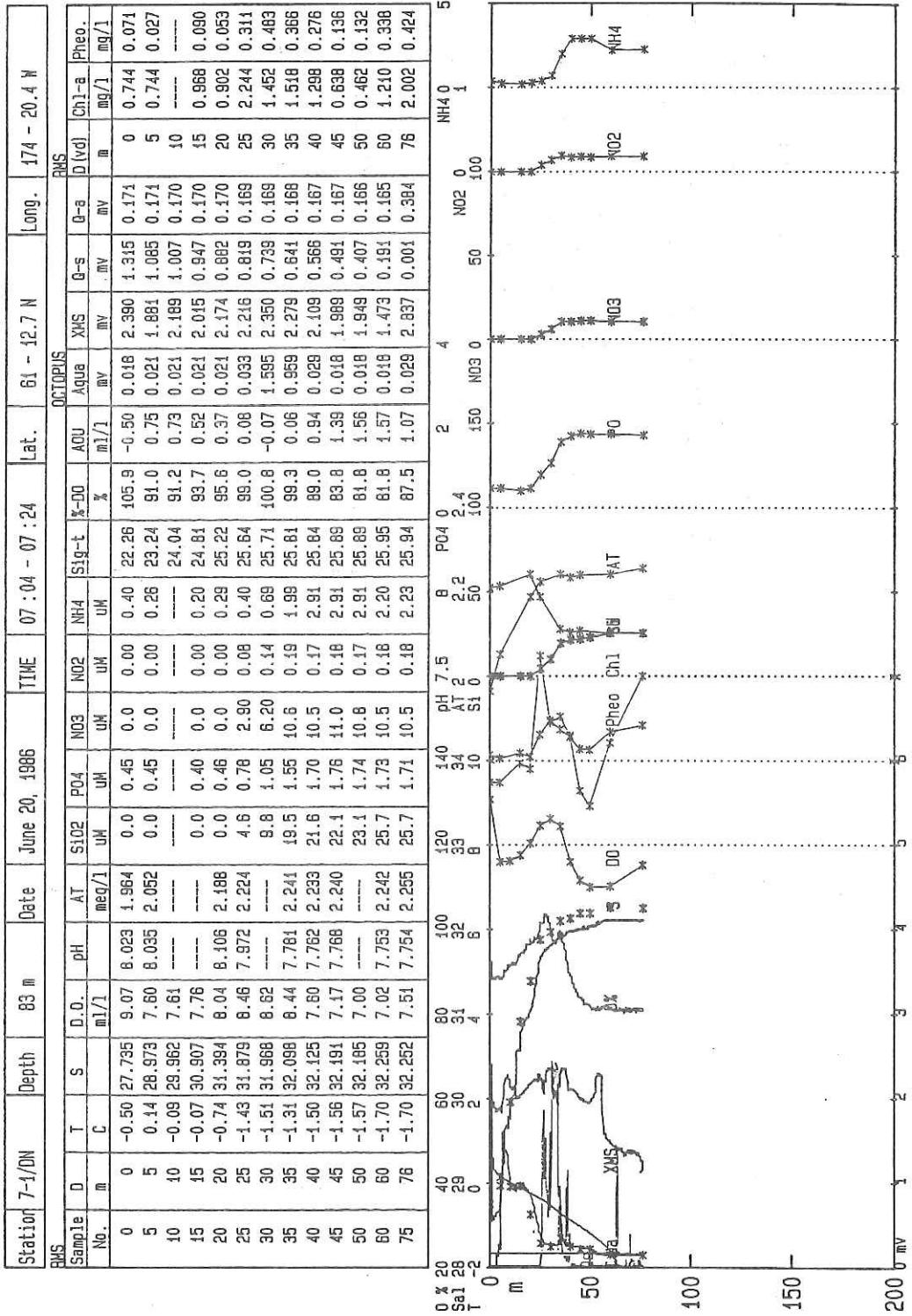


Table 30. Summary of OCTOPUS data at Station 7-2

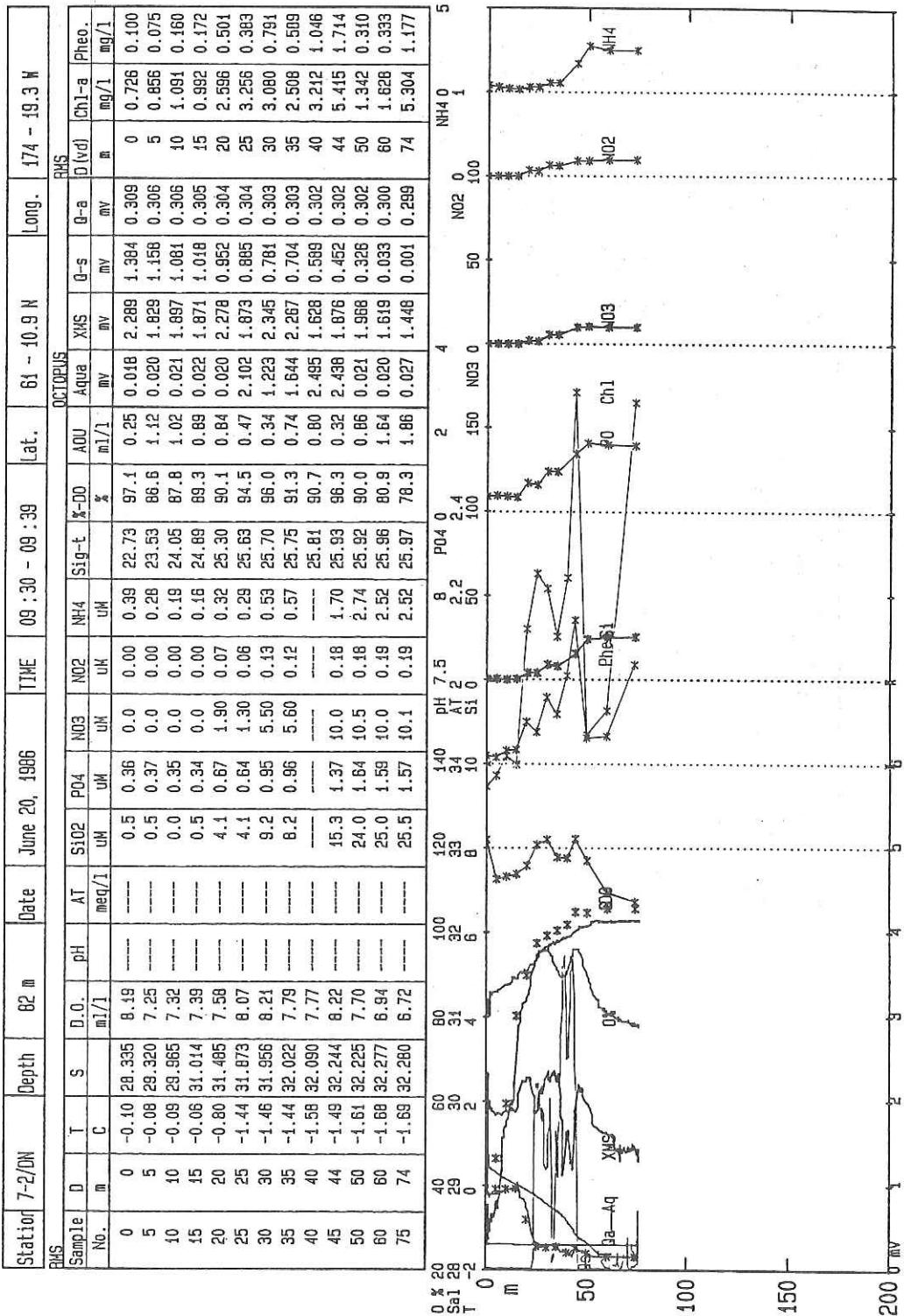


Table 31. Summary of OCTOPUS data at Station 7-3

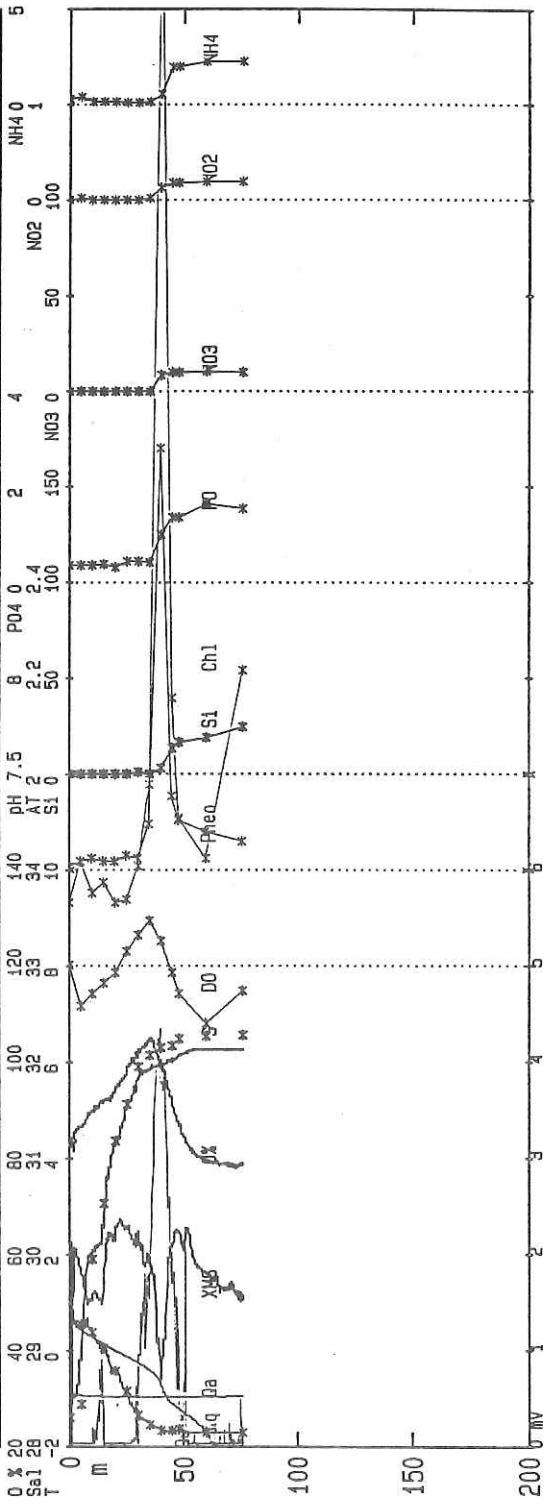


Table 32 Summary of OCTOPUS data at Station 7-4

RMS Sample No.	D m	T C	S ml/l	D.O. mg/l	pH	AT mg/l	SiO ₂		PO ₄		NO ₃		NH ₄		Sig-t		%±0		AOU		XHS		θ-S		θ-a		Dlyd		Chl-a		Pheo.	
							UW	UN	UW	UN	UW	UN	UW	UN	UW	UN	UW	UN	mv	mv	mv	mv	mv	mv	mv	mv	mg/l	mg/l				
0	0	2.20	28.485	7.08	8.015	2.004	0.5	0.30	0.0	0.0	0.29	22.74	89.2	0.86	0.032	0.863	1.453	0.531	0	0.645	0.062											
5	5	0.85	29.290	6.79	8.028	2.066	0.0	0.31	0.0	0.0	0.24	23.46	83.4	1.38	0.035	1.881	4.233	0.597	5	0.967	0.169											
10	10	1.00	29.954	7.05	8.019	2.099	0.0	0.30	0.0	0.0	0.21	23.99	87.0	1.06	0.035	1.608	1.162	0.539	10	1.203	0.150											
15	15	0.02	30.917	7.18	-----	-----	0.0	0.33	0.0	0.0	0.20	24.81	86.6	1.10	0.245	1.312	1.094	0.542	15	1.606	0.278											
20	20	-0.14	31.355	7.27	8.044	2.177	0.0	0.29	0.0	0.0	0.16	25.17	87.9	1.01	1.318	1.732	1.015	0.546	20	1.012	0.175											
25	25	-0.36	31.568	7.57	8.056	2.208	1.5	0.35	0.0	0.0	0.12	25.35	91.1	0.74	0.094	2.361	0.940	0.549	25	0.886	0.248											
30	30	-1.30	31.710	7.76	-----	-----	2.4	0.35	0.0	0.0	0.12	25.50	91.1	0.76	0.090	2.655	0.879	0.554	30	0.792	0.240											
35	35	-1.44	31.852	7.84	8.061	2.216	2.8	0.34	0.0	0.0	0.13	25.61	91.8	0.70	1.055	2.566	0.818	0.556	35	1.342	0.464											
40	40	-1.56	31.972	8.08	7.982	2.224	8.7	0.61	3.80	0.08	0.25	25.71	94.4	0.48	2.410	2.322	0.755	0.563	40	1.540	0.783											
45	45	-1.29	32.049	7.66	7.805	2.231	24.2	1.32	11.3	0.18	1.94	25.77	90.2	0.83	0.232	2.521	0.676	0.568	45	0.660	0.372											
50	50	-1.19	32.162	7.14	-----	-----	23.7	1.20	11.0	0.19	1.52	25.86	84.4	1.32	0.031	2.648	0.816	0.573	50	0.248	0.217											
60	60	-1.65	32.271	6.67	7.768	2.240	24.7	1.43	11.4	0.15	2.65	25.96	77.9	1.89	0.032	2.302	0.505	0.583	60	0.781	0.397											
75	75	-1.69	32.319	6.72	7.766	2.250	24.7	1.47	11.5	0.15	2.67	26.00	78.3	1.86	0.035	2.169	0.248	0.604	75	1.870	0.324											

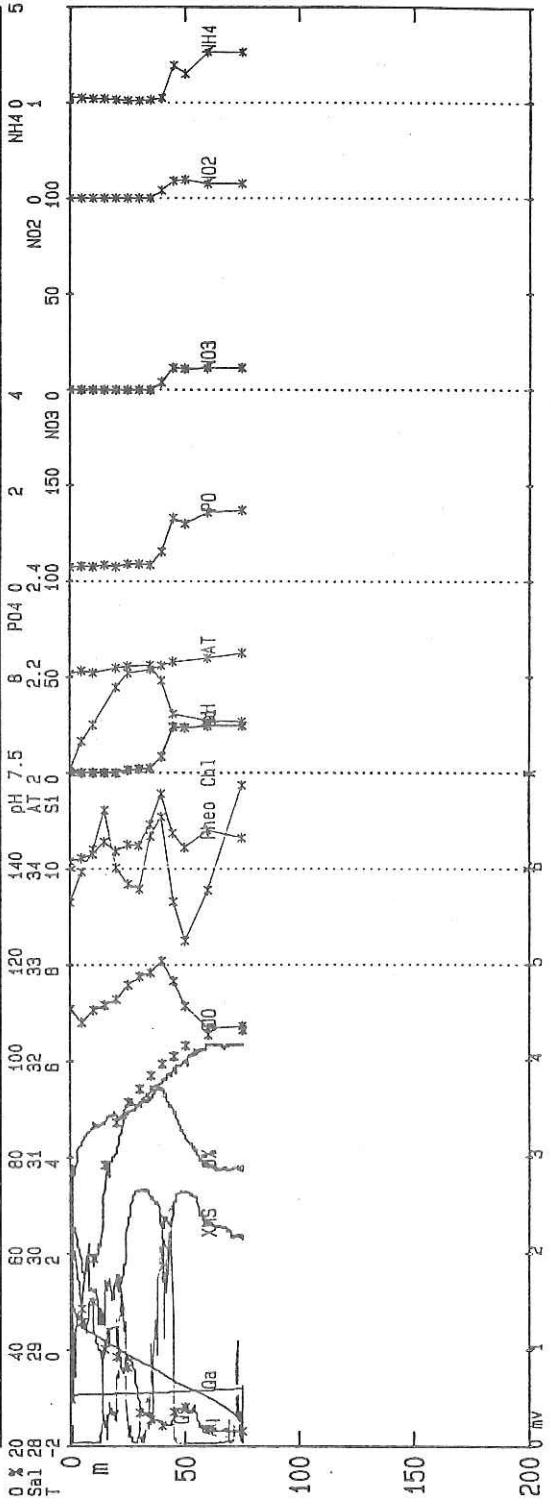


Table 33. Summary of OCTOPUS data at Station 8

Station	B/DN	Depth	110 m	Date	June 21, 1986	TIME	15:55 - 16:07	Lat.	56° 30.7' N	Long.	167° 47.2' W	
Sample No.	D m	T C	S	D.O. ml/l	pH	AT mEq/l	S102 um	PO4 um	NO3 um	NH4 um	Sig-t %	AOU ml/l
0	0	5.86	34.720	7.45	8.130	2.228	0.0	0.31	0.0	0.00	0.28	24.98
5	5	5.84	34.732	6.17	-----	-----	0.0	0.30	0.0	0.00	0.21	24.99
15	15	5.82	34.724	6.21	8.132	2.226	0.0	0.32	0.0	0.00	0.23	24.99
25	25	5.70	34.748	6.26	8.132	2.228	0.0	0.32	0.0	0.00	0.33	25.02
35	34	4.41	34.846	6.42	8.119	2.233	0.0	0.48	0.0	0.00	1.55	25.24
45	45	3.36	31.895	6.41	-----	-----	11.0	0.96	3.00	0.05	4.55	25.38
55	55	2.45	31.948	6.27	7.983	2.224	9.7	1.01	4.20	0.08	4.94	25.49
65	65	3.30	32.150	6.25	7.980	2.232	21.0	1.13	5.10	0.10	5.88	25.58
75	75	3.14	32.475	5.46	7.700	2.231	38.9	2.05	22.2	0.16	2.15	25.68
85	84	3.19	32.486	4.95	-----	45.6	2.22	24.5	0.46	2.20	25.86	25.77
95	95	3.21	32.497	4.77	7.642	2.239	46.1	2.23	24.5	0.48	2.43	25.87
105	106	3.23	32.501	4.66	7.639	2.242	46.1	2.31	23.0	0.48	2.13	25.87

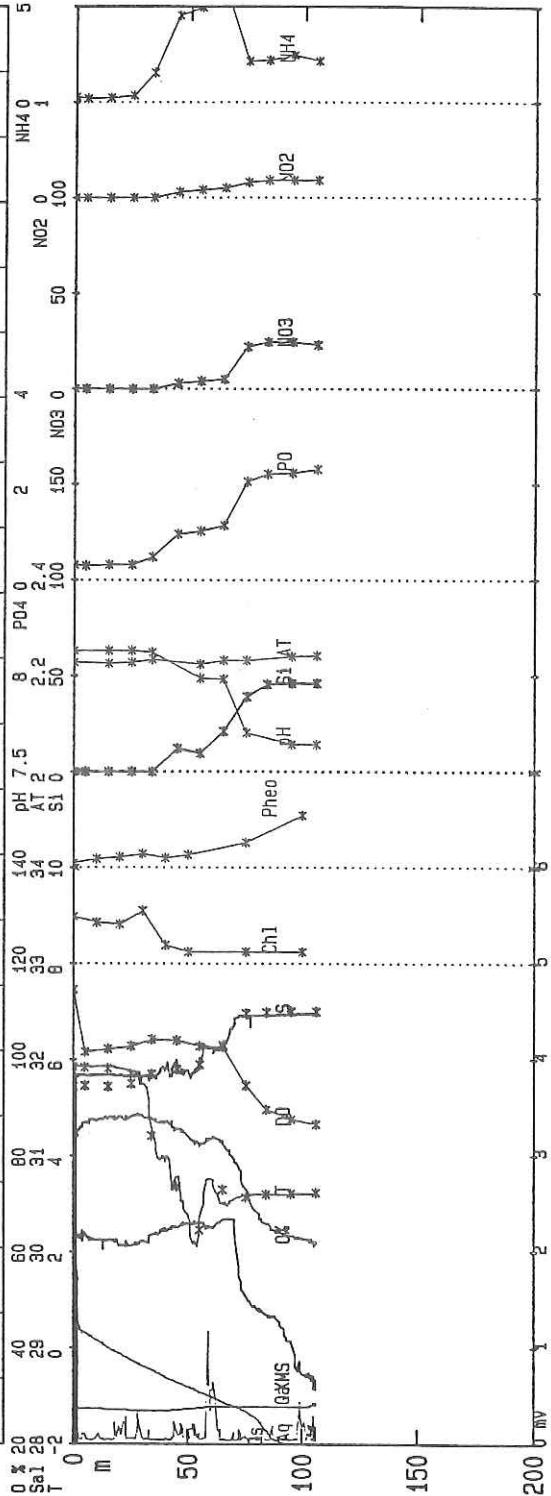


Table 34. Summary of OCTOPUS data at Station F'

Station	F'/DN	Depth	4105 m	Date	July 15, 1986	TIME	10 : 48 - 11 : 10		Lat.		53 - 00.2 N		Long.		144 - 59.2 W					
							D	T	S	O.O.	pH	AT	SI02	P04	NO3	NO2	NH4	Sig-t	%-DO	RHS
No.	m	C					meg/l	um	um	um	um		um	um	um	um	um	mv	mg/l	
0	0	0	8.86	32.205	7.37	----	13.3	1.12	10.0	0.07	0.27	24.95	111.7	-0.77	2.548	0.842	1.512	0.890	0	0.434 0.049
15	15	15	8.79	32.221	6.08	----	13.4	0.58	10.1	0.13	0.22	24.98	92.1	0.52	2.117	2.336	1.218	0.914	15	0.368 0.049
30	30	30	8.58	32.224	6.25	----	15.2	1.22	11.7	0.14	0.24	25.44	92.9	0.50	2.343	2.434	1.094	0.932	30	----
40	41	41	6.37	32.387	6.48	----	15.9	1.24	12.2	0.16	0.15	25.46	92.4	0.53	2.355	2.418	0.894	0.925	49	0.340 0.178
50	49	49	6.25	32.396	6.47	----	15.9	1.29	12.3	0.16	0.35	25.54	90.4	0.68	2.261	2.565	0.785	0.912	60	0.292 0.187
60	60	60	5.90	32.433	6.37	----	17.1	1.40	13.9	0.34	0.41	25.72	88.7	0.81	2.054	2.651	0.702	0.902	70	0.287 0.195
70	70	70	4.98	32.529	6.39	----	20.7	1.52	16.9	0.18	0.03	25.63	86.6	0.98	1.840	2.715	0.633	0.891	79	0.194 0.119
80	79	79	4.62	32.625	6.29	----	26.2	1.63	20.0	0.03	0.05	26.00	82.9	1.24	1.603	2.698	0.555	0.884	90	0.154 0.109
90	90	90	4.39	32.798	6.05	----	37.8	2.02	25.8	0.00	0.05	26.26	77.0	1.68	1.401	2.746	0.487	0.860	99	0.098 0.060
100	99	99	4.17	33.101	5.64	----	46.3	2.27	30.4	0.01	0.08	26.45	67.9	2.35	1.385	2.794	0.407	0.879	110	0.043 0.049
110	110	110	4.04	33.331	4.98	----	56.7	2.55	34.7	0.00	0.05	26.63	59.8	2.98	1.304	2.795	0.325	0.875	120	0.019 0.036
125	120	120	3.89	33.529	4.39	----	63.4	2.66	37.1	0.00	0.05	26.74	51.9	3.54	1.404	2.743	0.222	0.871	130	0.027 0.209
130	130	130	3.70	33.654	3.82	----														

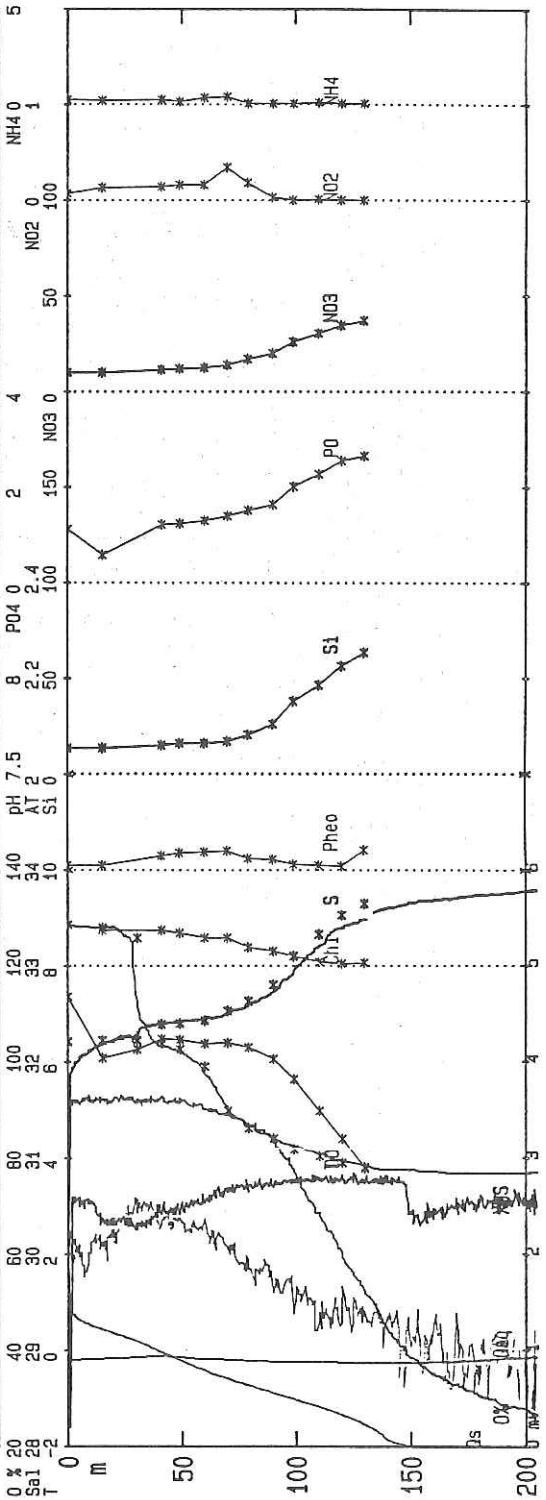


Table 35. Summary of OCTOPUS data at Station C'

Station	C°/DN	Depth	3920 m	Date	July 22, 1986	TIME	11 : 47 - 12 : 10			Lat.	53 - 30.3 N	Long.	177 - 31.2 E											
							S	D.0.	ph	AT	S102	P04	N03	NH4	N02	UW	UW	UW	%DO	ADU	Aqua	XHS	O-s	O-a
No.	m	C					med/1	UW	UW	UW	UW	UW	UW	UW	UW	UW	UW	UW	ml/1	ml/1	ml/1	ml/1	ml/1	mg/1
0	0	7.61	32.680	7.42	---	---	16.5	1.42	15.9	0.48	0.38	25.51	109.6	-0.65	0.636	0.913	1.409	0.425	0	0.533	0.450			
10	10	7.48	32.755	6.04	---	---	17.9	1.43	16.1	0.48	0.52	25.59	69.1	0.74	0.786	1.428	1.128	0.430	10	0.533	0.465			
20	20	7.41	32.755	6.04	---	---	17.9	1.34	16.1	0.48	0.58	25.59	89.0	0.75	2.162	1.416	1.025	0.438	20	0.608	0.163			
30	30	6.92	32.851	6.07	---	---	18.5	1.32	16.1	0.48	0.53	25.74	88.4	0.80	2.418	1.460	0.94	0.447	24	0.595	0.205			
40	40	6.71	32.851	6.17	---	---	19.7	1.20	16.7	0.20	0.53	25.76	89.3	0.74	2.383	1.602	0.787	0.456	29	0.570	0.230			
50	50	6.02	32.904	6.15	---	---	21.5	1.44	17.7	0.24	0.80	25.89	87.7	0.66	2.262	1.692	0.666	0.457	39	0.843	0.335			
60	60	4.74	33.050	6.32	---	---	25.7	1.60	19.8	0.29	0.66	26.16	87.4	0.91	2.068	1.809	0.553	0.455	49	0.744	0.361			
70	70	3.97	33.089	6.27	---	---	33.5	1.72	23.3	0.50	0.37	26.27	85.1	1.10	1.616	1.918	0.441	0.452	59	0.570	0.273			
80	80	3.77	33.121	6.13	---	---	36.5	1.78	24.8	0.21	0.37	26.31	82.9	1.26	0.737	2.039	0.338	0.446	78	0.434	0.337			
90	90	3.50	33.159	5.97	---	---	38.3	1.89	25.7	0.05	0.03	26.37	80.3	1.47	0.493	2.081	0.223	0.437	98	0.113	0.147			
100	100	3.34	33.166	5.87	---	---	40.7	1.91	26.9	0.01	0.48	26.39	78.6	1.60	0.247	2.081	0.088	0.449	146	0.023	0.063			
150	150	3.62	33.437	4.58	---	---	59.2	2.18	33.2	0.01	0.11	26.58	61.8	2.83	0.055	1.902	0.001	0.444	195	0.011	0.065			
200	200	3.91	33.745	3.17	---	---	74.8	2.56	38.7	0.02	0.15	26.77	43.2	4.17	0.104	1.695	0.001	0.398						

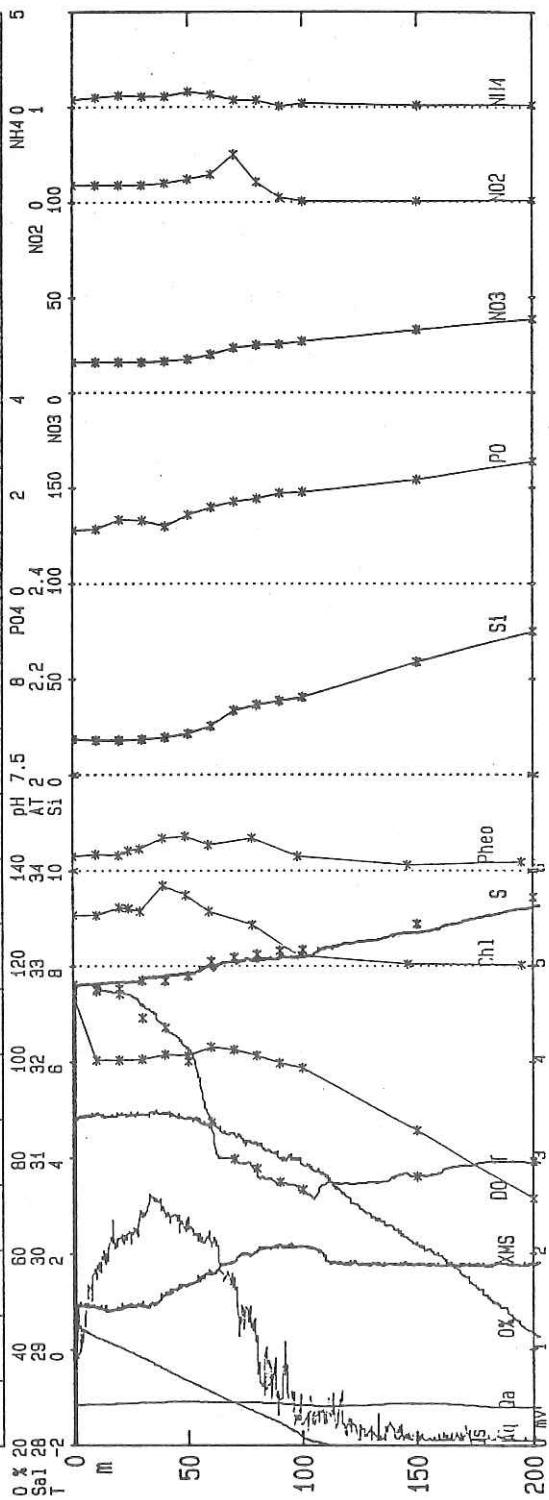


Table 36. Vertical distribution of total dissolved nitrogen (μg atoms N/l)

Sta.B	Sta.C		Sta.1		Sta.3		Sta.0		Sta.8		Sta.E		Sta.F		Sta.G		Sta.F'		Sta.C'		Sta.H	
	m	m	m	m	m	m	m	m	m	m	m	m	m	m	m	m	m	m	m	m	m	m
0	34.9	0	27.9	0	7.8	0	7.2	0	10.3	0	21.3	0	19.4	0	22.3	0	23.5	0	23.6	0	23.8	
15	34.8	15	26.6	10	6.7	5	7.5	15	9.4	10	22.2	10	24.3	10	20.3	10	21.7	10	24.2	10	23.8	
20	33.7	20	26.2	20	8.6	10	8.1	15	8.3	20	23.3	20	23.5	20	20.6	20	20.9	20	24.6	20	25.1	
25	34.9	25	29.2	40	28.1	15	9.5	25	7.3	25	22.9	25	18.3	25	20.1	25	24.9	25	27.4			
30	37.4	30	29.7	60	30.3	20	9.5	35	8.7	30	26.6	30	23.5	30	17.5	30	21.1	30	25.5	30	30.6	
40	35.9	40	32.3	60	30.1	25	11.3	45	16.6	40	27.2	40	21.5	40	17.7	40	19.3	40	26.4	40	31.5	
50	35.7	50	32.3	120	31.0	30	12.3	55	15.6	50	26.5	50	21.5	50	19.5	50	20.2	50	26.7	50	33.6	
60	35.7	60	34.0	65	34.0	35	13.3	65	18.8	60	23.3	60	18.8	60	19.6	60	28.9	60	35.3			
80	36.0	80	33.8	80	33.8	40	23.5	75	31.6	80	28.7	80	24.1	80	19.5	80	21.5	80	32.7	80	35.7	
100	37.1	100	38.4	95	34.0	85	32.9	100	34.0	100	25.1	100	21.5	100	26.0	100	33.8	100	38.6			
150	50.3	150	39.5	105	33.1	150	50.2	150	47.7	150	33.0	150	44.9	150	39.2	150	41.9	150	46.5	200	46.5	
200	56.1	200	44.2	200	32.1	200	51.9	200	50.5	200	40.3	200	48.6	200	45.6	200	45.6	200	46.5			
0	34.0	0	27.8	0	21.4	0	21.3	0	20.7	0	19.3	0	20.2	0	23.9	0	-	0	-	50	34.3	
50	33.5	50	32.2	10	26.5	50	24.2	50	21.0	50	20.3	50	19.3	50	26.3	50	-	50	-			
100	35.7	100	28.3	20	30.8	100	33.9	100	26.9	100	23.1	100	24.3	100	34.9	100	39.4					
200	55.4	200	44.3	30	31.7	200	52.0	200	50.2	200	41.4	200	48.0	200	45.6	200	46.7					
300	54.6	300	48.5	40	32.9	300	53.3	300	52.3	300	46.5	300	51.0	300	50.6	300	50.6					
400	54.3	400	51.6	50	32.5	400	52.1	400	50.9	400	50.9	400	50.9	400	52.5	400	52.3					
500	55.0	500	51.1	75	35.1	500	52.0	500	50.7	500	50.2	500	50.2	500	51.8	500	51.8					
600	56.1	600	50.3	100	34.7	600	51.5	600	50.5	600	49.1	600	52.0	600	51.3	600	53.3					
700	54.6	700	51.8	200	43.0	700	53.0	700	51.2	700	49.5	700	51.9	700	50.4	700	51.4					
800	55.4	800	52.1	300	48.4	800	51.8	800	52.2	800	51.5	800	51.5	800	50.1	800	52.1					
900	55.9	900	50.8	500	51.3	900	53.2	900	52.9	900	49.4	900	51.7	900	52.3	900	52.6					
1000	54.2	1000	51.3	750	51.4	1000	51.8	1000	51.3	1000	51.3	1000	54.4	1000	52.0	1000	52.3					
1250	55.4	1250	50.8	1000	51.0	1250	53.3	1250	50.4	1250	49.7	1250	50.7	1250	51.0	1250	51.9					
1500	55.4	1500	50.8	1500	50.8	1500	52.5	1500	48.6	1500	50.8	1500	52.5	1500	53.7							
1750	52.1	1750	51.1	1750	48.4	1750	51.9	1750	48.4	1750	49.3	1750	49.8	1750	50.5	1750	50.7					
2000	50.7	2000	47.8	2000	47.9	2000	47.9	2000	49.3	2000	49.0	2000	49.0	2000	50.6	2000	50.5					
2250	48.8	2250	47.0	2250	47.4	2250	48.6	2250	47.5	2250	47.5	2250	46.4	2250	50.7							
2500	47.7	2500	45.6	2500	45.6	2500	50.0	2500	48.1	2500	47.3	2500	47.7	2500	50.6							
2750	47.1	2750	44.7	2750	49.4	2750	47.7	2750	48.6	2750	47.0	2750	46.6	2750	49.8							
3000	48.8	3000	44.8	3000	48.8	3000	45.0	3000	48.0	3000	45.2	3000	44.6	3000	48.5							
3250	48.4	3250	43.9	3250	49.1	3250	47.7	3250	44.8	3250	44.8	3250	44.6	3250	48.9							
3500	45.2	3500	42.5	3500	46.6	3500	45.1	3500	47.1	3500	42.8	3500	45.0	3500	48.5							
3750	45.9	3750	43.4	3750	45.7	3750	43.5	3750	46.7	3750	44.6	3750	44.6	3750	48.4							
4000	45.4	4000	44.3	4000	45.4	4000	45.4	4000	45.4	4000	45.4	4000	45.4	4000	48.7							

* Water samples collected by Niskin bottle casts (Stas. C, F, G, F', G', C', F and H) or OCTOPUS casts (Stas. 3, D and 8) from shallow depths (upper portion of the table) and by CTD casts from a whole water column (lower portion of the table) were used.

Table 37. Vertical distribution of particle number (per ml)
and particle volume ($\mu\text{m}^3/\text{ml} \times 1000$)

Depth m	PN	PV	Depth m	PN	PV	Depth m	PN	PV	Depth m	PN	PV	Depth m	PN	PV
Sta.B (Nis)*			Sta.C (Nis)			Sta.1 (Nis)			Sta.2 (OCT)			Sta.3 (VD)		
0	1783	547	0	1725	459	0	3241	1003	0	1820	398	0	2772	974
15	1980	1366	15	2085	430	10	1761	998	10	2373	934	10	1806	851
20	1951	834	20	2310	396	20	1664	981	20	2403	689	20	3780	707
25	1616	492	25	1507	237	30	737	972	40	504	340	30	11836	2395
30	1304	531	30	1212	339	40	609	120	50	434	323	40	8420	1160
40	712	138	40	572	60	50	490	88	70	331	122	50	3317	194
50	473	196	50	719	97	75	424	118	80	415	49	75	2231	428
60	441	218	60	520	119	100	336	82	90	392	56	100	1091	220
80	332	34	80	446	195	200	303	217	110	577	88	140	3894	659
100	492	282	100	435	75				125	1107	24			
150	397	94	150	398	103									
200	103	10	200	491	52									
Sta.4 (Nis)			Sta.D (VD)			Sta.7-4 (VD)			Sta.8 (VD)			Sta.E (VD)		
0	3344	1502	0	865	864	0	7832	784	0	967	141	0	1027	353
			10	1330	2051	5	7231	771	10	1157	884	10	1121	702
			20	1207	4906	10	5967	734	20	1026	587	20	1128	731
Sta.5			30	1695	1591	20	5561	2438	30	1031	475	25	1181	451
			40	9301	4667	30	4238	566	40	1247	432	30	1355	622
0	1587	278				50	1874	325	50	1701	594	40	783	298
						70	3401	1320	75	1937	498	50	650	748
						80	4746	548	100	3740	1412	60	539	188
Sta.6												80	420	541
0	1387	990										100	350	105
												150	195	77
												200	124	118
Sta.F (Nis)			Sta.G (Nis)			Sta.F' (Nis)			Sta.C' (Nis)			Sta.H (Nis)		
0	926	98	0	1501	999	0	1643	197	0	2499	1093	0	3686	1472
10	871	83	10	1393	148	10	1513	610	10	2519	1091	10	3972	1502
20	755	95	20	1251	131	20	1204	362	20	2165	522	20	3068	556
25	648	314	25	1371	413	25	1055	220	25	2227	957	25	2821	562
30	514	66	30	1838	871	30	1309	691	30	2122	709	30	1971	523
40	538	154	40	1783	275	40	1295	168	40	1976	725	40	836	363
50	410	339	50	1575	692	50	1221	385	50	1579	226	50	741	4356
60	345	84	60	893	180	60	772	558	60	1239	282	60	614	62
80	262	146	80	386	351	80	428	124	80	1127	557	80	522	369
100	190	94	100	177	139	100	277	44	100	447	190	100	409	141
150	219	23	150	171	149	150	301	22	150	410	82	150	494	50
200	117	8	200	237	998	200	217	128	200	1132	115	200	856	131

* Water samples collected by Niskin (Nis), Van Dorn (VD) or OCTOPUS (OCT) casts were used.

Hydrographic characteristics

T. Nakai, H. Otobe and A. Hattori

1. The northwestern North Pacific and the deep Bering Sea (Stations B, H, C and 1)

On the western side of the Emperor Sea Mounts in the northwestern North Pacific (Station B), a cold watermass with a temperature less than 2.5°C lay between 60 and 150m, and formed a dichothermal layer below where a strong halocline was found. Mesothermal water was located at a depth of about 350m. These patterns are typical of the Western Subarctic Water (Fig. 11, see also Fig. 18).

On the eastern side of the Emperor Sea Mounts, south of the Near Islands (Station H), the distribution of temperature and salinity differed either from those at Station B or those at Station C located in the Bower's Basin of the Bering Sea. The dichothermal layer was hardly identifiable (Fig. 17) and mesothermal water higher than 4°C was seen at ca. 150m. The influence of the Alaskan Stream Water is apparent in this area.

At Station C in the Bower's Basin, the dichothermal layer was thick. This watermass probably originates from mode water with a temperature of approximately 2.2°C , salinity of 33.25 ppt and dissolved oxygen of 6.8 ml/l; the mode water was clearly seen at a depth between 100 and 150 m at Station 1 (Fig. 18). The inflow of the Bering Slope Current along the continental slope is suggested to be the possible source.

The concentration of dissolved silicic acid in the bottom layer of Station C was very high ($> 200 \mu\text{M}$) (Fig. 13), as observed in the other areas of the deep Bering Sea.

2. Northwestern Bering Sea shelf region (Stations 2-6 and D)

In the northwestern shelf region, the water structure was complex (Fig. 11), because of the seasonal cooling and heating, the vertical and horizontal water mixing by the tidal current and the wind, the freezing and melting of ice and runoff from large rivers. The temperature was very low. We encountered large floating ices at the vicinity of Station 4 on July 18-20. The surface salinity was very low (about 31.0 ppt), and cold water less than -1°C was commonly found near the bottom of the continental shelf area north of Station 4. The maximum oxygen content

exceeding 9 ml/l existed at about 30 m (Fig. 12). Silicic acid and nitrate in the surface layer were completely depleted (Figs. 13 and 14). Maximum concentration of chlorophyll *a* was found in the subsurface layer (30-40m), and the concentration of ammonium was very high in the bottom layer. It is suggested that the spring bloom of phytoplankton occurred before we made the observations.

3. The Gulf of Alaska (Stations E, F and G, Figs. 15 and 16)

Three stations 300 nautical miles apart (E, F and G) were occupied along 145°W meridian. The contours of sigma-t generally inclined toward the south, suggesting the occurrence of eastward geostrophic flow. The surface geostrophic velocity relative to 3000 db and the volume transport were calculated to be 1.1 cm/sec and $3.2 \times 10^6 \text{ m}^3/\text{sec}$ between Stations E and F, respectively, and 3.6 cm/sec and $6.8 \times 10^6 \text{ m}^3/\text{sec}$ between Stations F and G, respectively.

At Station F, the thermocline was at about 50 m. The dichothermal water was relatively warm and was hardly distinguishable from the mesothermal water (Fig. 17, top). The halocline and oxycline were located between 100 and 150 m (Fig. 17, middle and bottom). Judging from these profiles, the watermass at Station F can be identified as the Eastern Subarctic Water.

4. Water circulation in the studied area

The Alaskan Stream which originates from the Eastern Subarctic Water flows westward on the southern side of the Aleutian Islands. A great part of the stream enters into the Bering Sea through the Near Strait and the other part passes around the Near Islands together with a small part of the Western Subarctic water. The surface water flows eastward on the northern side of the Aleutian Islands, turns northwest in a direction near the shelf break area (the Bering Slope Current), and produces a cyclonic circulation over the deep Bering Sea basin. A part of the circulating water flows along the Kamchatka Peninsula (the East Kamchatka Current) and serves as the source of the Western Subarctic Water. A part of the East Kamchatka Current is drifted eastward by the westerly wind (the Subarctic Current), while the other part contributes to the formation of the Oyashio Current. The T-S and T-O₂ diagrams illustrated on the basis of the CTDO data collected in this cruise (Fig. 18) are consistent with this general pattern of water circulation. The

data on surface currents estimated from the ship's drift (Fig. 19) also generally support this view, but some discrepancy can be noted. This may be attributed to the temporal variations of the flow pattern and the presence of vortex currents.

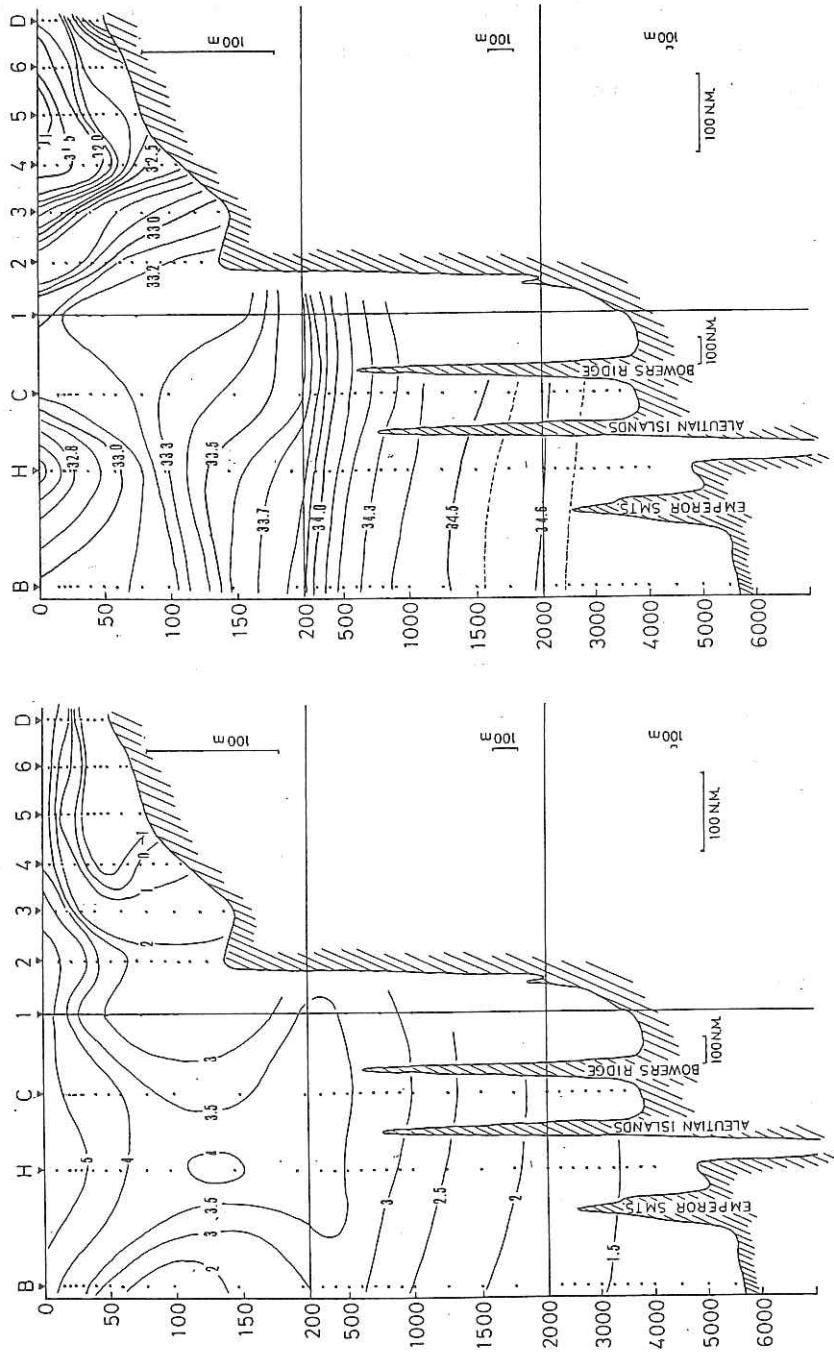


Fig. 11. Cross sections of temperature (C) and salinity (ppt) along a transect between Station B and D.

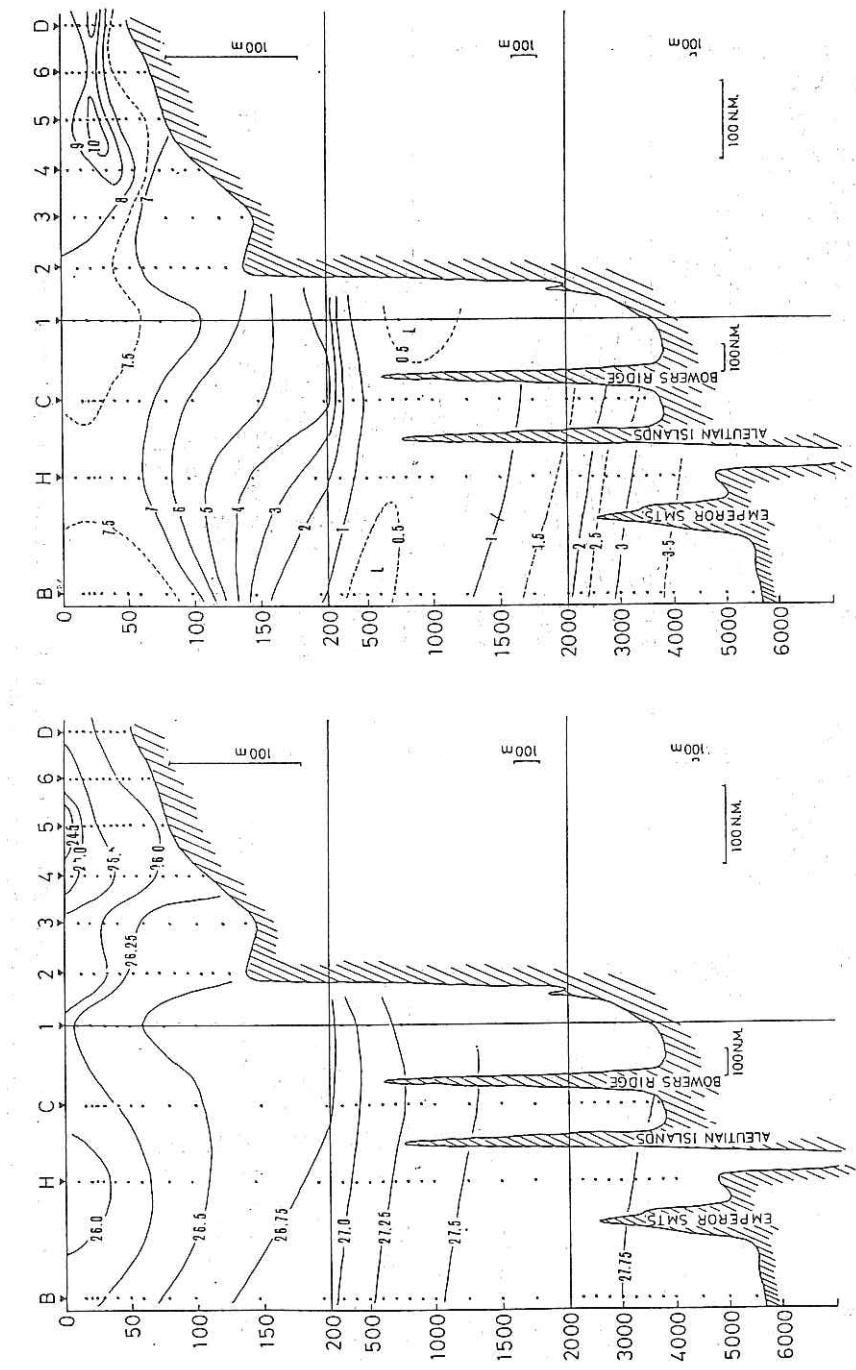


Fig. 12. Cross sections of sigma-t and dissolved oxygen (ml/l) along a transect between Stations B and D.

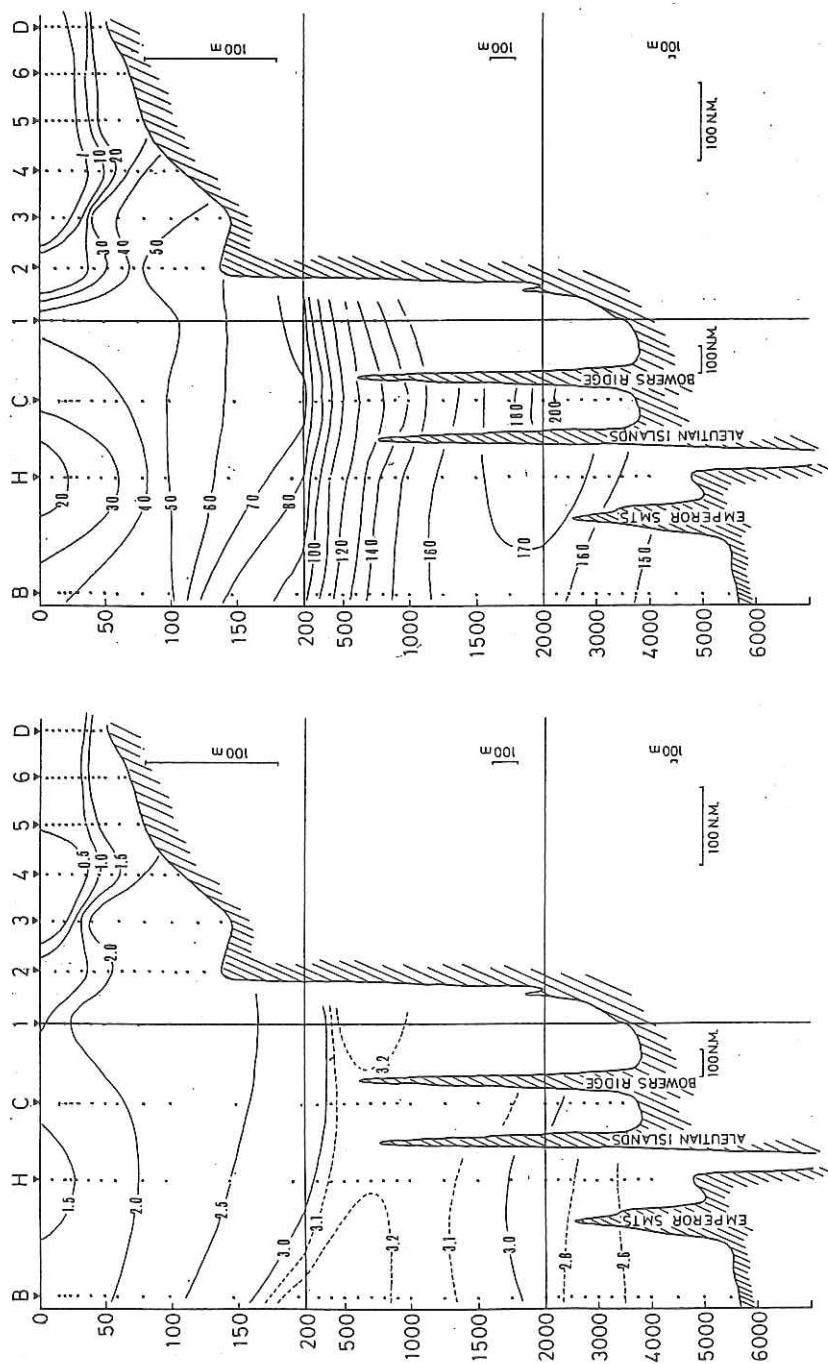


Fig. 13. Cross sections of phosphate ($\mu\text{g atoms P}/1$) and silicic acid ($\mu\text{g atoms Si}/1$) along a transect between Stations B and D.

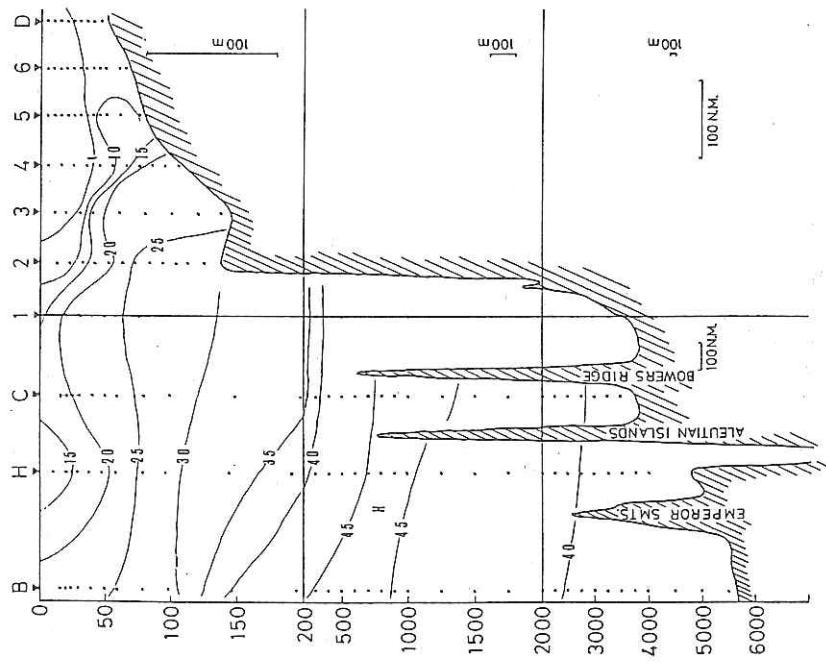


Fig. 14. Cross sections of nitrate ($\mu\text{g atoms N/l}$) along a transect between Stations B and D.

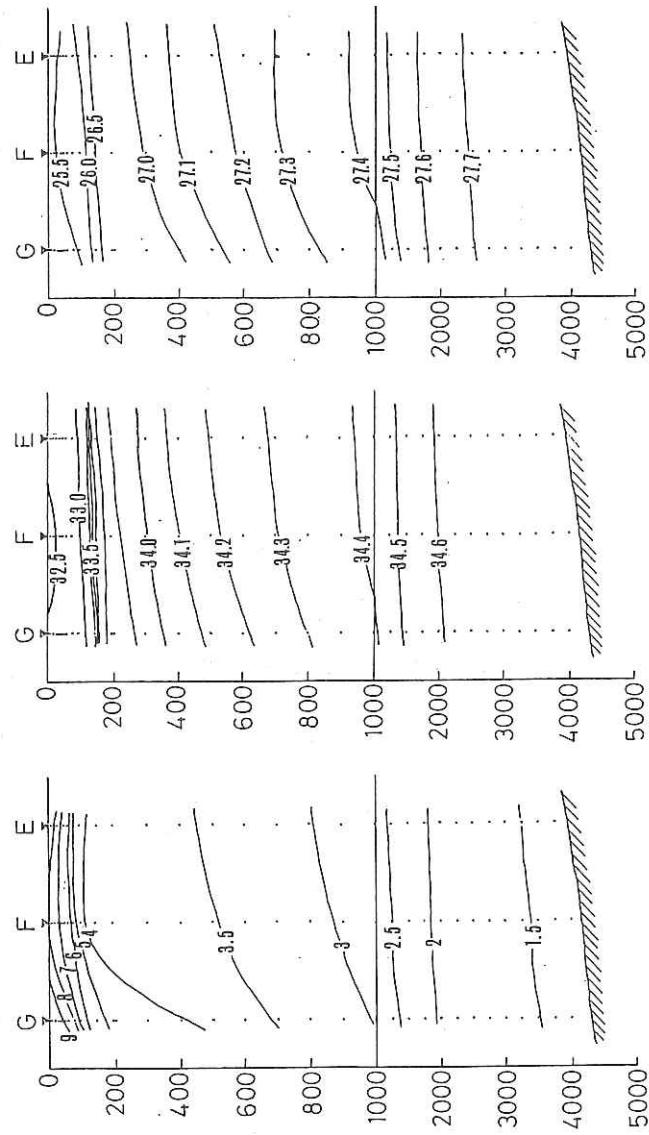


Fig. 15. Cross sections of temperature (C), salinity (ppt) and sigma-t along a north-south transect in the Gulf of Alaska.

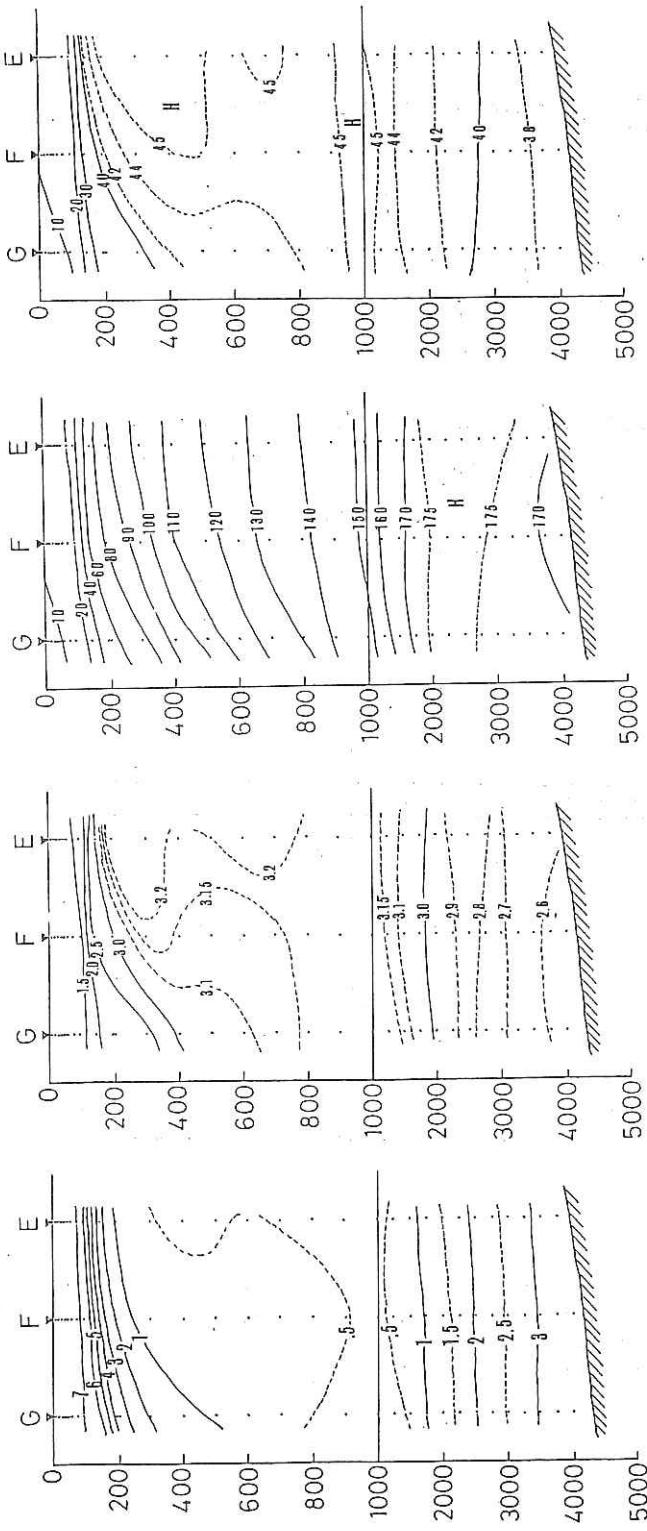


Fig. 16. Cross sections of dissolved oxygen (ml/l), phosphate ($\mu\text{g atoms P/l}$), silicic acid ($\mu\text{g atoms Si/l}$) and nitrate ($\mu\text{g atoms N/l}$) along a north-south transect in the Gulf of Alaska.

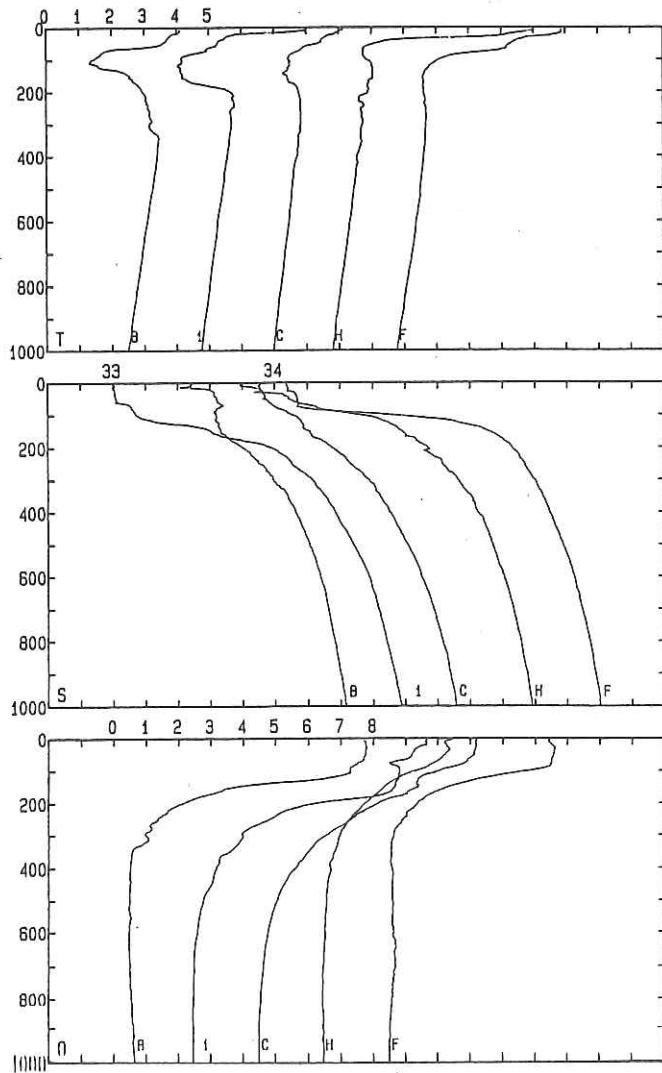


Fig. 17. CTDO profiles of temperature (C, top), salinity (ppt, middle) and dissolved oxygen (ml/l, bottom) at Station B, I, C, H and F. For illustration, origins of abscissa scale for the individual station data were successively shifted toward the right by two units.

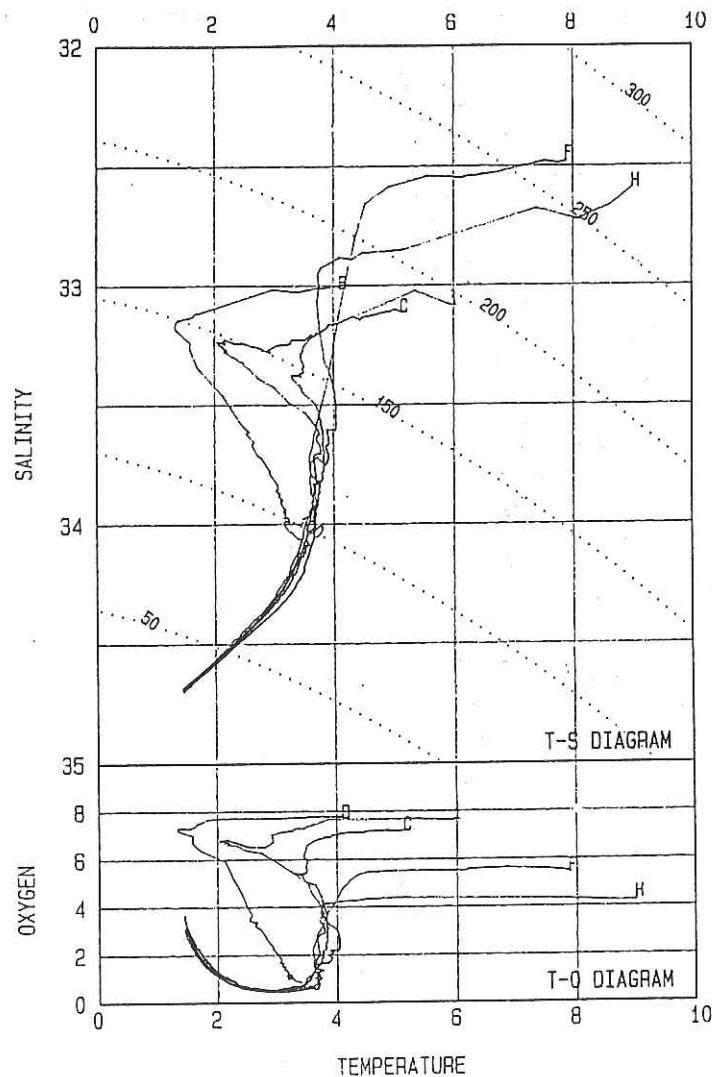


Fig. 18. T-S and T-O₂ diagrams at Stations B, 1, C, H and F. Data collected by CTDO casts were used.

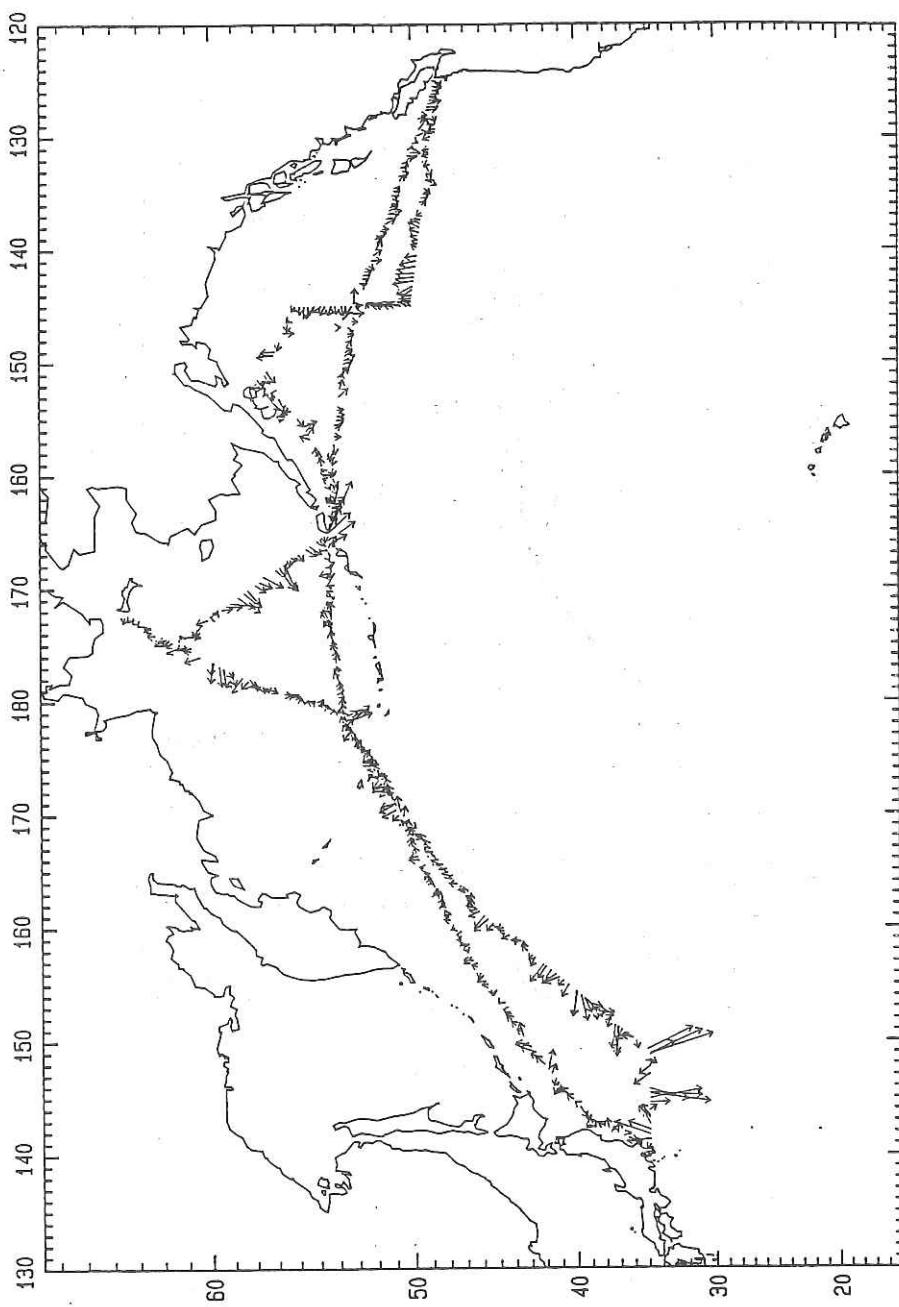


Fig. 19. Surface currents estimated from the ship's drift.

Water temperature observation in the upper ocean

H. Otobe, T. Saino, K. Harada and H. Sasaki

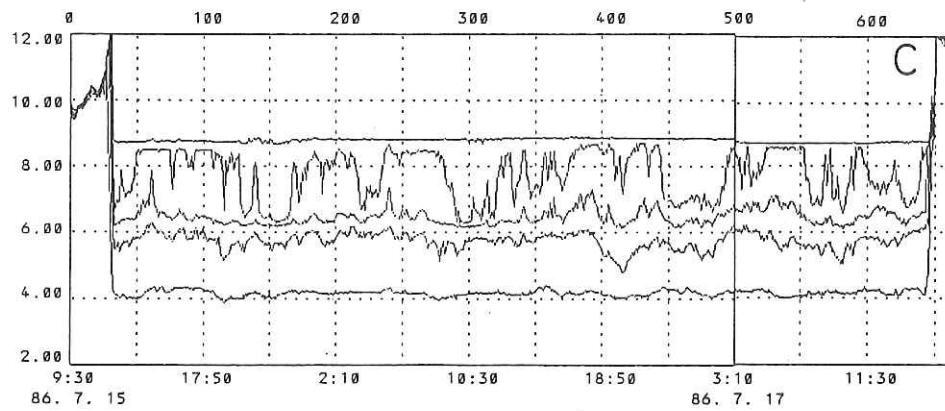
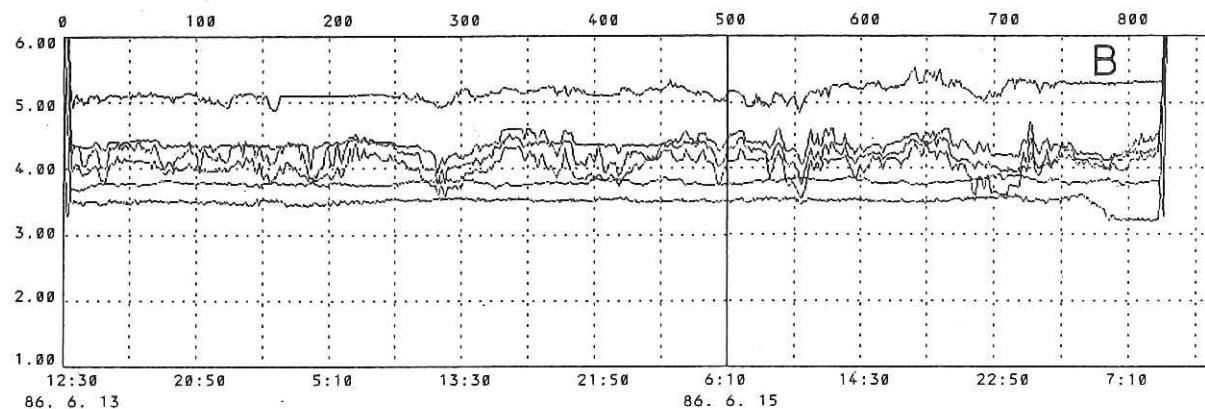
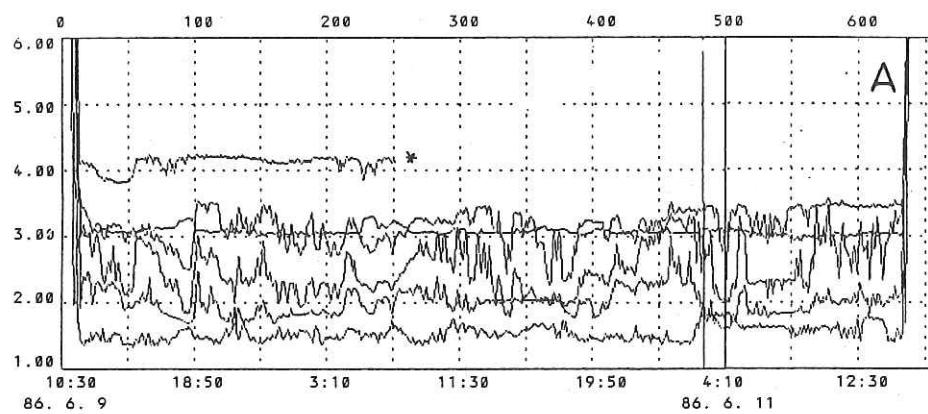
In order to obtain information on the advective water movement, water temperature in the upper layer was continuously monitored in the Bering Sea and the Gulf of Alaska. Six self-contained digital thermometers (Model RMT water thermometer, Rigosha & Co., LTD, Tokyo) with an accuracy of $\pm 0.05^{\circ}\text{C}$ and resolving power of 0.02°C were attached to a sediment trap buoy system of the Ocean Research Institute, University of Tokyo and 2 thermometers to that of Hokkaido University (Table 38).

The data stored in a memory unit of each thermometer (4096) were read out on a personal computer through a RC-232C serial interface and transferred to mini floppy disks.

The records are reproduced in Fig. 20.

Table 38. Summary of water temperature observations

Station	Observation period	Sampling interval	Observation depth intended	Buoy system used	
				min	m
B	1100 9 Jun-1530 11 Jun	5	10,40,50,60,100,200		ORI
C	1300 13 Jun-0900 16 Jun		10,40,50,60,100,200	(floating)	
F'	1150 15 Jul-1530 17 Jul		15,30,40,60,100		
C	1200 13 Jun-0900 22 Jul	15	100,250		Hokkaido Univ.
F	0800 29 Jun-0900 15 Jul	10	260,600		(anchored)



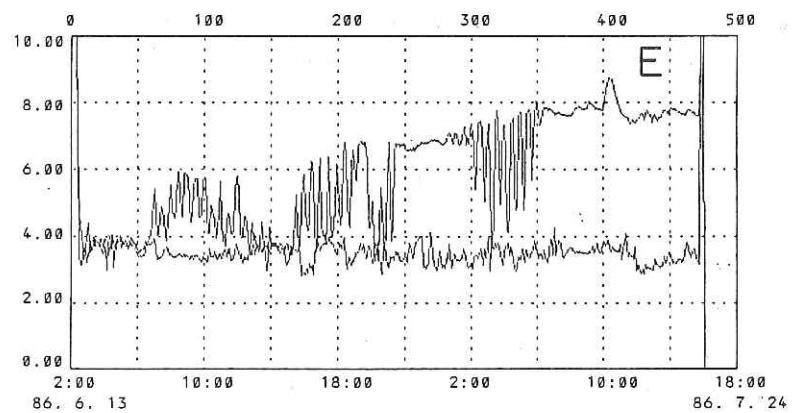
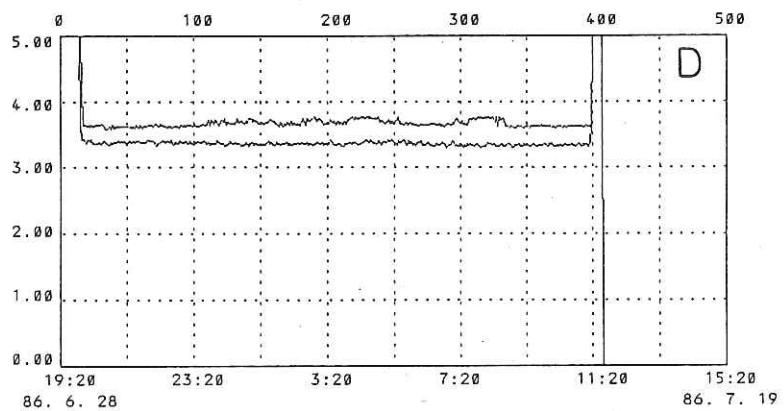


Fig. 20. Water temperature records at 10, 40, 50, 60, 100 and 200 m (from top to bottom) at Station B during the period from 1100 LST 09 June to 1530 LST 11 June 1986 (A), at Station C during the period from 1300 LST 13 June to 0900 LST 16 June 1986 (B), at 15, 30, 40, 60 and 100 m at Station F' during the period from 1150 LST 15 July to 1530 LST 17 July 1986 (C), at ca. 100 and 250 m at Station B during the period during 1200 LST 13 June and 0900 LST 22 July 1986 (D), and at depths of 10-60 m and 350-400 m at Station F during the period from 0800 LST 29 June to 0900 LST 15 July 1986 (E).

Chemical and radiochemical studies of the northern
North Pacific and the Bering Sea

S. Tsunogai, K. Harada, K. Taguchi, S. Nagao and K. Demura

1. Sediment trap experiments

We collected settling particles with two types of sediment traps (D and NH types) at three stations and studied the spatial, temporal and vertical variations of particulate fluxes with special reference to the high biological productivity in this region. Emphasis was placed on fluxes of chemical and radiochemical components. One of the three arrays of time-series sediment traps (D type) which had been deployed at Station B on 11 August 1985 during the KH-85-4 cruise was retrieved on 9 June 1986, and the other two arrays of sediment traps were installed from 13 June to 22 July 1986 at Station C in the Bering Sea Basin and from 29 June to 15 July 1986 at Station F in the Gulf of Alaska, respectively (Fig. 21). The particle samples were collected on Nuclepore filters (pore size, 0.6 μm) and freeze-dried. Total mass fluxes are listed in Table 39. Major chemical components, trace metals and natural radionuclides in these samples will be determined.

2. Radionuclides in seawater

In order to study the removal processes of reactive chemical elements from the ocean, the following natural radionuclides in seawater were determined: ^{234}Th , ^{232}Th , ^{230}Th , ^{228}Th , ^{231}Pa , ^{210}Po , ^{210}Bi and ^{210}Pb .

The samples for ^{234}Th , ^{210}Pb , ^{210}Bi and ^{210}Po analysis (20-50 l each) were collected from 10 depths in the upper 100 m layer at Stations C and F' with a submersible pump and also collected from 11 depths in the bottom 500 m layer at Station B, C and F' with Niskin bottles. On the basis of the results obtained, the effects of biological activity and the bottom interface on the removal of these nuclides from seawater will be investigated.

The samples for the determination of long lived radionuclides, ^{232}Th , ^{230}Th , ^{228}Th and ^{231}Pa (180-200 l each), were collected from 16 and 17 depths at Stations C and F', respectively, with large volume water samplers and a submersible pump. Surface samples (about 200 l each) were also collected at 26 stations, and ^{228}Th content was determined. The results will give information for understanding the slow removal

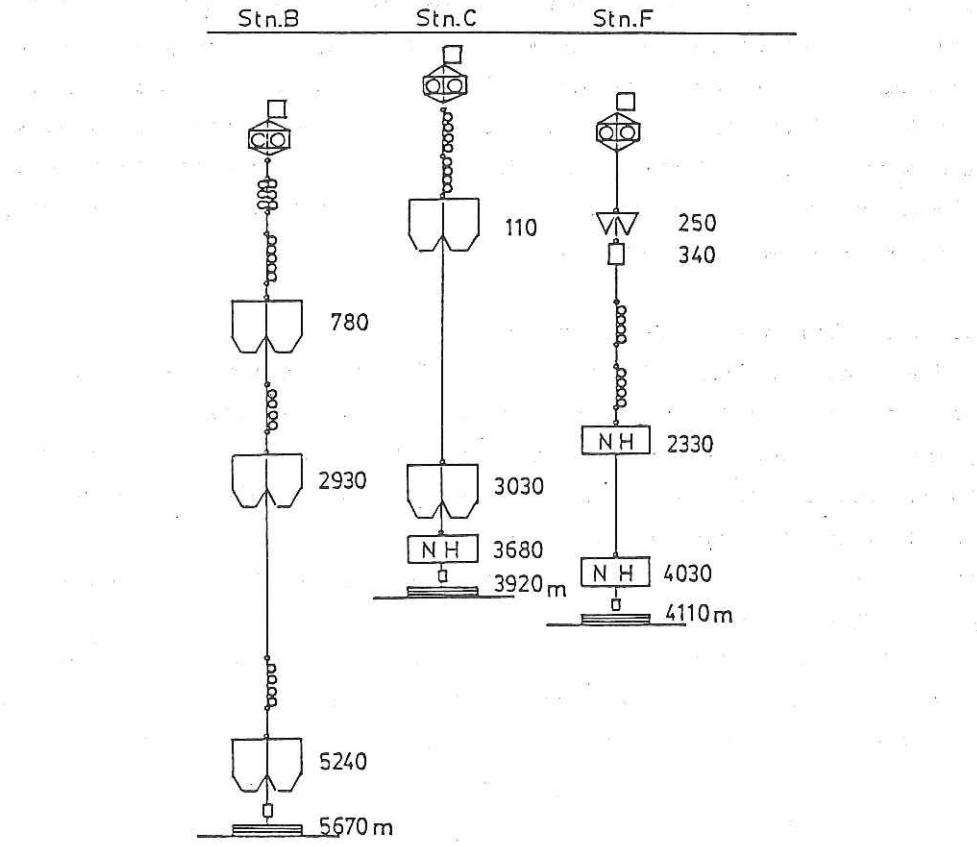


Fig. 21. Arrays of sediment traps used at Stations B, C and F.

processes of these elements in the ocean.

3. Uranium and some heavy metals in marine sediments

Sediment samples were obtained with a large box corer at Stations B, C and 3 and with an Okean grab sampler at Station F'. The core samples were cut into 5 cm segments, and interstitial water was separated using a squeezer made of polyvinylchloride under mechanical pressure. The interstitial water samples (about 500 ml each) will be analyzed for uranium and other heavy metals. The solid fractions of the sediment samples will be also analyzed for uranium and other metals after water and organic matter contents are determined. The data can be used to elucidate the behavior of uranium and heavy metals during the early diagenesis of marine sediments.

4. Dimethylsulfide in the ocean water

The distribution of dimethylsulfide (DMS) in seawater was studied in order to estimate the emission rate of DMS from the ocean to the atmosphere. The samples for the DMS measurements (about 50 ml each) were taken from the surface 200 m layer at 17 stations with Niskin bottles and analyzed for DMS and other reduced sulfur compounds on board the vessel within 12 hours by FPD-gas chromatography. Some of the results are given in Table 40.

At Stations C and F', we obtained 18 bottom water samples to disclose a sink term of DMS in the ocean. The results given in Table 41 show that the concentrations of DMS near the sea-floor are negligibly small.

5. ^{210}Pb in maritime aerosols

Aerosols in maritime air were collected by filtration through a Whatman 41 filter (20 x 25 cm) using a large volume air sampler(Fuji VFC 304P, flow rate 1 m^3/min). The air sampler was set on the compass bridge deck of the vessel and a sector controller was used to avoid contamination from the ship exhaust.

Twenty one samples were obtained during the cruise. The atmospheric concentrations of ^{210}Pb were determined by digesting the filter samples with hot nitric acid and by using a radiochemical technique devised by Tsunogai and Kondo (1982). The beta activity of ^{210}Pb was counted with a low back-ground gas flow type GM counter.

Table 39. Total particulate fluxes observed with sediment traps

Type of trap	Depth m	Period					Total mass flux	
							mg/m ² •day	
Station B (46°49'N, 162°07'E, water depth 5670 m)								
D	780	11 Aug '85-24 Oct '85	74 days	309	634* ¹			
		24 Oct '85- 6 Jan '86	74	105	194* ¹			
		6 Jan '85-21 Mar '86	74	65	107* ¹			
		21 Mar '85- 3 Jun '86	74	111	229* ¹			
D	2930	11 Aug '85- 9 Jun '86	302	170	179* ¹			
D	5240	11 Aug '85-24 Oct '85	74	258	329* ¹			
		24 Oct '85- 6 Jan '86	74	51	91* ¹			
		6 Jan '85-21 Mar '85	74	33	47* ¹			
		21 Mar '85- 3 Jun '85	74	55	102* ¹			
Station C (53°30'N, 177°31'E, water depth 3920 m)								
D	110	13 Jun '86-22 Jun '86	9	21	142* ²			
		22 Jun '86- 1 Jul '86	9	26	137* ²			
		1 Jul '86-10 Jul '85	9	24	40* ²			
		10 Jul '86-19 Jul '85	9	24	126* ²			
D	3030	13 Jun '86-22 Jun '86	9	10	3* ²			
		22 Jun '86- 1 Jul '86	9	295	912* ²			
		1 Jul '86-10 Jul '86	9	120	86* ²			
		10 Jul '86-19 Jul '86	9	97	84* ²			
NH	3680	13 Jun '86-22 Jul '86	39	478 ± 63 (n=6)				
Station F (53°00'N, 145°00'W, water depth 4110 m)								
S* ³	250	29 Jun '86-15 Jul '86	16	73				
O* ⁴	340	29 Jun '86-15 Jul '86	16	66				
NH	2330	29 Jun '86-15 Jul '86	16	127 ± 36 (n=5)				
NH	4030	29 Jun '86-15 Jul '86	16	121 ± 43 (n=5)				

^{*¹ Formaldehyde was used as a preservative.}^{*² Sodium azide was used as a preservative.}^{*³ A funnel-shaped trap with bubbles and without a lid (ORI-UT).}^{*⁴ A cylinder-shaped trap without a lid (Tohoku Univ.).}

Table 40. Dimethylsulfide concentrations in seawater near the surface

Station B		Station C		Station D		Station F	
Depth (m)	DMS (ng/l)	Depth (m)	DMS (ng/l)	Depth (m)	DMS (ng/l)	Depth (m)	DMS (ng/l)
0	95	0	31	0	6	0	2.0
15	56	15	20	5	15	10	4.2
20	74	20	40	10	20	20	5.4
24	135	30	47	20	22	24	5.7
29	197	40	37	25	20	29	2.6
39	80	50	54	30	16	39	8.9
59	23	59	19	41	8.4	49	7.9
78	21	79	23	45	16	59	4.2
98	6.9	99	10			78	10
195	5.8					98	3.4
						146	6.7
						195	0.6

Table 41. Dimethylsulfide concentrations in the bottom waters*

Station C (water depth 3920 m)		Station F' (water depth 4070 m)	
Depth (m)	DMS (ng/l)	Depth (m)	DMS (ng/l)
3620	1.8	3849	0
3720	0.8	3784	0
3821	0.8	3875	0
3846	0.5	3923	0
3870	0.5	3971	0
3890	0.5	4019	0
3899	0	4048	0
3909	0	4057	0
3914	0	4061	0

* Detection limit is 0.2 ng DMS/l.

Vertical distribution of total dissolved nitrogen in the
northern North Pacific Ocean and the Bering Sea

I. Koike and L. Tupas

The vertical distribution of total dissolved nitrogen was determined for stations B, C, E, F, and G in the northern North Pacific Ocean and for stations 1, 3, D, and 8 in the Bering Sea. Water samples from shallow depths (surface to 200 m) were collected with 23-liter Niskin bottles, while deep water samples were collected with a rosette multi-sampler.

Water samples were first filtered through Whatman GF/F filters, stored at 4°C and were analyzed on board within 10 hours after collection. Total dissolved nitrogen was determined with a Yanaco Total Nitrogen Analyzer (Model TN-7). Forty μ l of sea water was injected manually into the reaction column of the machine and 4 to 6 trials per water sample were performed. Water samples were preserved after analysis with mercury chloride.

Preliminary results (Table 36) show that waters of the northern North Pacific Ocean contain between 17.5 to 56.1 μ g atoms N/l of dissolved total nitrogen while waters of the Bering Sea contain between 21.4 to 51.4 μ g atoms N/l at the deep basin (maximum depth sampled at 1000 m) and between 7.2 to 33.1 μ g atoms N/l at the shallower area (maximum depth sampled at 105 m).

CO₂ exchange between the air and sea

S. Kanamori, T. Aono, N. Kadoya and A. Hashimoto

The high productivity of the Bering Sea has been considered to be supported by active input of the deep water rich in nutrients from the Pacific. We can expect a close relation between carbonate and nutrient concentrations. Therefore, we investigated the dynamics of carbonate and nutrients in association with production and decomposition of organic matter, and CO₂ exchange between air and sea in this productive area.

The total dissolved carbonate and individual carbonate species were calculated from pH and titration alkalinity. The pH was measured, within a few hours after sampling with an air-tight pH cell system at $25.0 \pm 0.1^\circ\text{C}$ on the NBS standard buffer scale. The precision of the measurement was ± 0.003 pH unit and the reproducibility within ± 0.001 pH unit.

The titration of alkalinity was carried out by a computer processed automatic potentiometric titration system with an accuracy of $\pm 0.1\%$ error.

The measurements of pH was carried out at all stations and those of alkalinity at Stations B, C, 1, F, F', C', H (Tables 4-12).

Distribution of arsenic

S. Kanamori and A. Hashimoto

Arsenic is present in two oxidation states in seawater, arsenate and arsenite. The thermodynamically stable form is arsenate. Arsenite is probably produced by biological reduction of arsenate. The occurrence of organic derivatives of arsenic in natural waters has been reported. It is suggested that the dissolved states of arsenic in seawater are affected or controlled by biological activity. We investigated the distribution and dissolved states of arsenic species in the Bering Sea and the northern North Pacific where the primary productivity and other biological activities are known to be very high.

Seawater samples were collected at Stations B, C, D, E and F. One liter of sample water was filtered through a Nuclepore filter ($0.4 \mu\text{m}$) within a few hours after the sampling. One hundred ml of the filtrate was kept frozen and the remaining filtrate was acidified with HCl. Chemical analysis will be made at Nagoya University.

Heavy metals in marine aerosols

S. Kanamori

Aerosol samples were collected for the study of the aerial and size distribution of heavy metal aerosols (Cu, Zn, Cd, Fe, Al, and Mn) in the maritime air over the northern North Pacific and the Bering Sea. Aerosol particles greater than $0.4 \mu\text{m}$ were separated into 8 size fractions and the remaining smaller fraction was caught on a back-up filter (Nuclepore Filter, $0.4 \mu\text{m}$) by the use of an Andersen low volume sampler.

Ten sets of aerosol samples were collected during this cruise. Chemical analysis will be made at Nagoya University.

"Clean" water sampling and heavy metal study

S. Kanamori, T. Aono, N. Kadoya and A. Hashimoto

The distribution and dissolved states of several heavy metal elements (Cu, Zn, Cd, Fe, Al and Mn) were investigated. Emphasis was placed upon the organic-metal association and the nutrients-dissolved metal relation.

New "clean" water samplers, one for surface water and the other for subsurface water, were designed and constructed at the metal workshop of Nagoya University and used in this cruise for the first time. Because of their complex structure, the clean water samplers reported previously were inconvenient in operating on board and often failed to function properly. Our samplers were designed to eliminate these defects. A special device was made to avoid any contamination from the sampler itself, the steel wire, and the ambient water during operation. One set for surface water and four sets for subsurface water were operated at a time.

Seventeen casts were carried out during the cruise and "clean" seawater samples were taken from 62 layers covering from the surface to 5500 m depth. Although a few samplers did not work well in the early phase of trial, the reliability of water sampling was increased by improved handling. There was no mal-functioning in the last 5 casts.

Setting of the "clean" water sampler and processing of the sample water were done in a clean room (class 100) on board the ship. The capacity of the samplers was 1.8 l. The suspended particulate matter was separated by centrifugation (10,000 rpm for 1 hr.) or by filtration through a Nuclepore filter (0.4 μm). The sample waters were then processed through a column of reversed phase absorption resin, and a majority of metal-organic compounds was separated. The chemical analysis will be made at Nagoya University.

Distribution of vanadium and molybdenum in seawater

N. Kadoya

Vanadium and molybdenum are biologically active metals and their distributions in the oceans may reflect the mode and extent of biological activity. We have found that vanadium is depleted in the surface water of the Southern Sea where the biological production is very high. This study attempts to collect information on the distributions of vanadium and molybdenum in the highly productive Bering Sea and northern North Pacific where available data are extremely limited.

Sea water samples were collected at all stations. Vanadium was determined with an autoanalyzer on board the ship within several hours after sampling by the catalytic oxidation method of Bindschedler using 4,4'-bis(dimethylamino) diphenylamine and potassium bromate. The concentration of vanadium tended to increase with depth: 20 to 30 nmol/l in the surface layer and 35 to 40 nmol/l in deep water. One hundred ml water sample was saved for molybdenum analysis in the shore laboratory.

Study on the distribution of Se species, Se(IV), Se(VI) and organic Se,
in seawater and on the presence in gaseous state of Se and some
other bio-essential elements in maritime air over
the Bering Sea and the northern North Pacific

T. Aono, Y. Nakaguchi and K. Hiraki

The main purpose of the present study is to improve our understanding of the mechanisms which control the chemical state of Se in seawater.

The concurrent occurrence of Se (IV), Se (VI) and organic Se in natural waters has been reported. The chemical state of Se is regulated by the oxidation and reduction potential of ambient seawater and sediment and also by the action of microorganisms. The present cruise covers the Bering Sea and the northern North Pacific where biological activity is known to be very high. We determined three Se species in seawater from these areas to obtain information how the distribution of Se species reflects the biological activity.

Our previous studies upon trace bio-essential elements in dust, mist and fume in the atmosphere have shown that a substantial portion of these elements exists in gaseous state. By applying a cold-trap method using liquid nitrogen, we collected gaseous substances in maritime air over the Bering Sea and the northern North Pacific, and Se and some other bio-essential elements were determined. Information on the distribution of these elements in gaseous state and its spatial and temporal variation is crucial for understanding the geochemical cycle of the trace bioelements.

Air sampling for the study of increasing atmospheric methane

T. Yoshinari

The concentration of atmospheric methane is increasing at about 2 % per year. This rate of increase is approximately correlated with that of the human population on the earth. While major sources of methane have been identified, their relative source strengths are yet to be clarified.

We intend to establish a data base for the abundance of ^{13}C , ^{14}C and ^2H in methane of air samples from various parts of the surface of the earth. Based on the information and the data of the isotope ratio for major sources we hope to establish the source strengths of major sources on the earth.

Six air samples have been collected during the trans-Pacific cruise between Vancouver, Canada and Tokyo, Japan. Each sample was contained in an aluminum cylinder in which about 2.5 m^3 air was filled by using a compressor. These will be used for analyses of the abundance of ^{13}C , ^{14}C and ^2H in methane as well as the concentrations of H_2 , CO and CH_4 in air.

Depth profiles of nitrous oxide in the northern
North Pacific Ocean and Bering Sea

A. Hattori and I. Koike

Seawater samples, collected from standard hydrocasts of 23-l Niskin bottles (above 200 m depth) and 5-l Niskin bottles (below 200 m depth) at Stations B, C, F, and H, were transferred directly into 100 ml serum bottles using a siphon. After allowing ca. 200 ml of water to overflow, the bottles were capped with rubber stoppers pierced with a hypodermic needles to avoid air bubbles. In order to terminate the biological reaction, 0.2 ml of 0.1 M HgCl_2 was injected through the rubber stopper. Dissolved nitrous oxide was determined by the gas chromatographic method of Cohen (1977).

Reference

Cohen, Y. (1977), Shipboard measurement of dissolved nitrous oxide in seawater by electron capture gas chromatography. *Anal. Chem.*, 49, 1238-1240.

A study of maritime aerosol over the northern
North Pacific Ocean and Bering Sea

A. Ono and M. Yamato

The importance of a deeper understanding as to how and to what extent man's activities influence chemical processes in the atmosphere has become apparent. Remote oceanic areas are suitable fields for studying the transformation of aerosol particles in the atmosphere and the transport processes of pollutants from the continents to the atmosphere because there are no local sources of pollutants. Our main concern in this study is to clarify the molecular state of submicron sulfate particles over the oceans.

Samplings of gases and aerosol particles in maritime air were undertaken over extended areas of the northern North Pacific and Bering Sea. To avoid contamination by ship exhaust, aerosol samplers were placed on the upwind side of the stack of ship, about 13 m above the sea surface. The samplers were automatically controlled so as to operate only when the wind speed exceeded 3.6 m/sec and when the wind direction was within 60 degrees from the forward direction of the ship. For chemical analysis of bulk components, aerosol particles were collected with a large volume Andersen sampler. For electron microscopic examination, aerosol particles were collected on electron microscope grids coated with a thin film of carbon, calcium, nitron and barium chloride by the combined use of a two-stage impactor and an electrostatic aerosol sampler. Sulfur dioxide, a precursor of sulfate in aerosol, was collected on a filter paper soaked in Na_2CO_3 solution after the removal of aerosol particles by passing through a fluoropore filter. The concentration of total aerosol particles was determined using an expansion-type portable nuclei counter.

Sulfate and nitrate particles were identified by the method of Bigg et al. (1974), and the method of Isawa and Ono (1977), respectively. The sulfuric acid component in the aerosol particles was examined by the calcium thin film technique of Ono et el. (1983). A preliminary examination suggests that sulfuric acid particles predominate in remote oceanic areas. Over the Bering Sea almost all particles reacted completely with the calcium thin film.

References

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Geochemical study of beryllium isotopes

M. Kusakabe

The ratio of cosmogenic nuclides ^{7}Be (half life 53.6 days) to ^{10}Be (half life 1.5 million years) can be used as a geochronometer for phenomena in air and surface water. In addition, the $^{10}\text{Be}/^{9}\text{Be}$ ratio (^{9}Be is stable) has been used recently in geochronology with a time scale of several million years or more. However, the behavior of the nuclides on the earth surface is not well understood due to the difficulty in measurement.

In order to clarify the behavior of the Be isotopes, the following experiments have been done.

1. Distribution of Be isotopes in surface water

About 200 l of surface water samples were collected approximately every 24 hours while the ship was steaming. Preconcentration and purification of Be were carried out partially on board the ship in collaboration with the Hokkaido University group who are interested in U-Th series nuclides. After extraction of U-Th series nuclides, the samples were brought back to the laboratory. The activities of ^{7}Be have been counted by using gamma ray spectrometry. The samples will be further purified and analysed for ^{10}Be by an accelerator.

Measurement of ^{9}Be using gas chromatography will be done on 500 ml surface water samples which were taken at the same locations as above.

2. Vertical distributions of Be isotopes

At three stations (Stations B, C and F), vertical profiles were taken for Be isotopes. Seawater samples (40-50 l) were collected at Station B for ^{7}Be and ^{10}Be from 12 depths with Niskin bottles. More detailed profiles were available with large amount of water (170-200 l) at Stations C and F' by using a submersible pump and a large volume water sampler. The activities of ^{7}Be have been counted for surface water samples 200 m and above. ^{10}Be will be measured for every sample. The waters for ^{9}Be (~500 ml) were taken from the routine hydrocasts.

3. Be isotopes in settling particulates

Sediment trap samples from three locations (Stations B, C and F) were

available for ^{7}Be measurement (see Tsunogai et al. in this volume for details of the traps). The activity of ^{7}Be was detected in the sample from 1100 m depth at Station B. Weak but significant amounts of ^{7}Be activity were also found in the samples which had been collected with the traps deployed at 120 m at Station C and at 250 m at Station F. The evaluation of total flux of ^{7}Be can be done with the knowledge of the total flux of particulates which is not available yet. ^{9}Be and ^{10}Be will be measured for all samples obtained including the samples from deep waters.

4. Be isotopes in aerosol

Eighteen aerosol samples were taken along the cruise track by passing more than 1500 m³ of air through Whatman #541 filter paper. It took at least 1 day to get enough volume depending on weather conditions. The filter papers were brought back to USC and have been analysed for ^{7}Be by gamma ray spectrometry. Concentrations of ^{10}Be and ^{26}Al will be measured by an accelerator.

Study on biochemical constituents of particulate matter in deep water

E. Tanoue

Biochemical components of particulate matter, phytoplankton, zooplankton and deep-sea sediments were examined. Emphasis was placed on the clarification of the origin and fate of particulate organic matter in deep water.

Approximately 40 l of seawater was collected with Niskin bottles from the surface to 3000 m depth at Stations B, C, E, G, F' and H. Particulate matter was collected by filtering ca. 20 l of seawater through a precombusted Whatman GF/F (47 mm in diameter) glass fiber filter. Four hundred to six hundred liters of seawater was collected from 8 water layers from the surface to 2000 m depth at Stations B, C and F'. Two hundred liters of the seawater samples was filtered through a Whatmann GF/F (150 mm in diameter) glass fiber filter under a hydraulic pressure of ca. 200 mmHg. Ten liters of the filtrate was adjusted to pH 2.0 with HCl and heneicosanoic acid was added as an inner standard. Dissolved lipids were extracted three times with chloroform.

Plankton samples were collected by a NORPAC twin net or an ORI net at Stations B, C, E and D, and sediment samples by a box corer at Stations B, C and 3. All the samples were stored frozen (-30°C) until later chemical analyses in the land laboratory.

Organic C and N were determined using a CHN analyzer. Total carbohydrate, amino acid and lipid were determined by colorimetric, fluorometric and wet oxidation methods, respectively. Lipid materials were separated into simple lipid, glycolipid and phospholipid fractions. Simple lipid fraction was further separated into hydrocarbon, free fatty acid and mono-, di- and triglyceride fractions by silica gel chromatography. Lipid fractions were quantified as fatty acid contents by gas liquid chromatography. Proteins were solubilized with sodium dodecyl-sulfate (SDS), and molecular weight distributions of proteins and their subunits were determined by a SDS-polyacrylamide gel electrophoresis. Carbohydrate was hydrolyzed and monosaccharide components were determined by gas liquid chromatography. Chemical form of carbohydrate was further characterized by combined use of staining with a specific fluorescing dye and enzymatic digestion.

Natural abundance of ^{15}N and ^{13}C in suspended and
settling particulate organic matter

T. Saino and A. Hattori

Data on the natural abundances of ^{15}N and ^{13}C in the particulate organic matter (POM) provides useful information for understanding the processes of POM production and decomposition. During this cruise, we collected settling materials by using sediment traps and the suspended particulate materials with Niskin bottles, and determined their isotopic ratios to obtain insight into the particle dynamics in the oceanic waters.

Water samples were collected with 23-liter Niskin bottles at Stations B, C, F, F', C' at depths from 0 to 4000 m. Closely spaced water samplings were also carried out in the shallow layer (<200 m) at those stations. Surface water samples were obtained with a bucket at each station and once a day from 18 July to 29 July while steaming.

Sediment trap experiments (Table 42) were conducted at Stations B, C and F'. Sediment traps were hung at 10, 20, 30, 40, 50, 60, 70, 80, 100, 125, 150, and 200 m at Stations B and C, and at 15, 30, 40, 50, 60, 70, 80, 100, 125, 150 m at Station F'. The water samples (ca. 60 liters for >300 m, ca. 20 liters for <200 m) and the sediment trap samples were filtered through glass fiber filters (Whatman GF/F, 47 mm diameter, precombusted at 450 C for 4 hr), and POM collected on the filters was kept frozen for later mass spectroscopic analysis.

Table 42. Summary of the sediment trap experiments

Station	Date	Location		
B	Initial: 11:37 9 June	46	40.3'N	162 01.9'E
	Final: 15:09 11 June	46	42.5'N	162 27.6'E
C	Initial: 12:30 13 June	53	29.8'N	177 35.7'E
	Final: 09:00 16 June	52	59.4'N	178 26.3'E
F'	Initial: 12:13 15 July	52	59.6'N	144 56.1'W
	Final: 15:09 17 July	52	56.0'N	144 41.9'W

Diel variations of downward particulate flux at a 60 m depth
observed in a time-series sediment trap experiment

H. Sasaki and S. Nishizawa

The temporal variation of downward particulate flux within a day in the coastal water was documented by Welschmeyer et al. (1984). The variation indicates the occurrence of the diel change of zooplankton grazing activity in the upper water column. In the open ocean, an observation has been made in the western North Pacific by Oyama et al. (in press) using sediment traps which were repeatedly deployed and retrieved at 6-hour intervals. They showed a marked diel variation of particulate flux in a 60 m layer, 20 m beneath the subsurface chlorophyll maximum layer. The maximum flux was observed sometime in the evening.

During the Cruise of KH-86-3, diel variations of downward fluxes were determined using a newly designed time-series sediment trap (0.196 cm^2 in mouth area) which can collect 12 consecutive samples of sinking particles. A surface-floating array consisted of a main buoy, a rader buoy, a radio buoy and a 200 m suspension rope (cf. page 84) onto which the trap was fastened at a 60 m depth.

The experiments were conducted at Stations B, C and F'. Unfortunately, we failed to retrieve the sediment trap at Station C. Location of sampling sites, suspension periods and collecting intervals are shown in Table 43.

References

- Oyama, Y., H. Sasaki and S. Nishizawa (in press). Diel and daily variations of downward particulate flux in the upper 200 m water column. Preliminary Report of the Hakuho Maru Cruise KH-85-3.
- Welschmeyer, N. A., A. E. Copping, M. Vernet and C. J. Lorenzen (1984). Diel fluctuation in zooplankton grazing rate as determined from the downward vertical flux of pheopigments. Mar. Biol. 83, 263-270.

Table 43. Location of sampling sites, suspension periods of
the trap array, and collecting intervals

Station	Location	Date	Suspension	Collecting
B	46 39.9'N 162 01.9'E	9-11 June 1986	48 hr	4 hr
F'	52 59.9'N 144 55.9'W	15-17 July 1986	48 hr	6 hr

Carbon and nitrogen uptake by natural populations of phytoplankton, and regeneration and oxidation of ammonium in shallow oceanic waters

A. Hattori and M. Otsu

Uptake rates of carbon dioxide, ammonium and nitrate by natural populations of phytoplankton in shallow waters of the northern North Pacific and the Bering Sea were measured by a ^{13}C - ^{15}N dual tracer technique. Water samples were collected with 23-liter Niskin bottles from depths of 100, 25 and 3.6 % light penetration, and from depths of the ammonium maximum and nitrite maximum (cf. Tables 20-35). Subsamples (2 liters) were transferred to polycarbonate bottles after passing through a 200 μm mesh screen to remove a majority of zooplankton, and incubated with 0.2 μM of $\text{NaH}^{13}\text{CO}_3$ (90 atom %) and 1-2 μM of K^{15}NO_3 (99 atom %) or 0.1-0.5 μM of $(^{15}\text{NH}_4)_2\text{SO}_4$ (99 atom %) under constant illumination from a bank of daylight fluorescent lamps at a light intensity of 270 μEinst per $\text{m}^2 \cdot \text{sec}$. For the subsurface samples, light intensities were reduced to simulate natural light conditions using metal screens. After 5 and 10 hours of incubation, the samples were filtered through Whatman glass fiber filters (type F, precumbusted at 450° C for 4-5 hours), and the filters and filtrates were saved for later analysis of ^{13}C and ^{15}N .

Using the filter samples, amounts of particulate organic carbon and nitrogen and ^{13}C and ^{15}N contents were determined with a Nichiden-Anelva TE-360 quadrupole mass spectrometer. The uptake rates were calculated from increase of ^{13}C and ^{15}N enrichments in particulate matter with time and initial ^{13}C enrichment in carbonate and ^{15}N enrichment in ammonium or nitrate.

Ammonium in the filtrates was separated by steam-distillation in the presence of excess MgO. Nitrate was reduced to nitrite by passing through a copper-cadmium column, and treated successively with aniline and β -naphthol. The azo-dye produced was extracted with n-hexane. The amounts of ^{15}N in the ammonium and azo-dye were determined, and the rates of ammonium regeneration and of ammonium oxidation were estimated from change of ^{15}N in ammonium and nitrate with time.

The mass spectrometric analysis has not yet been completed. The data available at present are entered in Table 44.

Table 44. Carbon and nitrogen uptake by natural populations of phytoplankton

Station	Depth	Amb. conc. ($\mu\text{g atoms N or C/l}$)			Specific rate (1/hr)			Uptake rate($\text{ng atoms/l}\cdot\text{hr}$)		
		NH ₄	NO ₃	NO ₂	ΣCO_2	POC	V _{amm}	V _c	ρ_{amm}	ρ_{nit}
B	0	0.31	20.3	0.25	2220	2.01	14.6	2.68*	3.65**	1.94*
	9	0.55	20.0	0.25	2230	2.07	15.0	4.03	5.56	1.90
	22	0.36	21.4	0.22	2240	2.08	14.1	2.55	2.35	1.09
	50	0.98	22.4	0.23	2260	1.65	12.4	0.18	0.13	0.12
	100	0.57	28.7	0.27	2270	0.52	6.3	0.26	0.65	0.69
C	0	0.37	19.3	0.17	2220	2.39	17.4	4.22	3.39	1.60
	8	0.37	19.6	0.17	2220	2.52	18.1	3.69	3.34	1.91
	34	0.49	20.6	0.17	2220	1.72	14.8	3.64	3.57	0.81
	60	0.99	22.7	0.21	2230	0.85	10.4	0.12	0.23	-0.12
	80	0.66	24.7	0.24	2260	0.53	9.4	0.23	0.51	-0.03
1	0	0.06	13.8	0.11	2170	4.32	28.3	1.33	0.98	2.19
	7	0.06	13.9	0.12	2180	4.29	28.5	1.31	1.07	2.65
	25	0.43	22.5	0.23	2210	2.13	19.8	1.71	1.54	0.20
	50	0.20	24.1	0.27	2250	0.83	10.2	0.23	0.15	-0.16
	100	0.02	25.9	0.37	2250	0.53	8.4	0.25	0.25	0.00
D	0	0.01	0.0	0.01	2080	1.79	17.0	2.47	2.38	1.70
	5	0.00	0.0	0.01	2090	1.83	16.9	2.39	2.51	2.38
	21	0.00	0.0	0.02	2110	2.33	18.3	2.60	2.39	2.48
	45	4.45	9.4	0.13	2260	2.55	18.4	0.42	0.87	0.47
										0.10

* Rate estimated from ^{15}N or ^{13}C enrichments at 0 and 5 hr.

** Rate estimated from ^{15}N enrichments at 5 and 10 hr.

Distribution of planktonic protista

S. Hara and E. Tanoue

The composition, standing stock and distribution of phototrophic and heterotrophic members of protista and monera in seawater were examined.

Seawater samples (1 liter each) were collected from the surface to 200 m with Niskin bottles at Stations B, C, E, F, G, F', C' and H. Samples were also collected from the surface and down to near the bottom with Van Dorn samplers at shallow stations. Deep water samples were collected with Niskin bottles at Stations B, C, E, F' and H.

Phototrophic and heterotrophic protista and phototrophic monera (blue green algae) were fixed with glutaraldehyde (1% final concentration). These organisms were collected on a Gelman membrane filter (25 mm in diameter and 0.2 μm in pore size). The filter was mounted on a slide glass with glycerine jelly, stored at -20°C and examined by epifluorescent microscopy. Heterotrophic monera (bacteria), fixed with glutaraldehyde (1%), were stained with a fluorescing dye (DAPI), collected on a Nuclepore filter (25 mm in diameter and 0.2 μm in pore size), stored at 4°C and examined by an epifluorescesce technique. The community structure of planktonic protista and monera at each station will be investigated. The other part of the seawater samples was divided and treated with glutaraldehyde (1 %), Lugol's solution (5 %) or formalin (5 %). Morphology and morphometry of phototrophic and heterotrophic protista will be examined by light and electron microscopies.

Studies on the processes of production at low trophic levels
and the organic substances in the northern Pacific Ocean
and the Bering Sea

K. Tada and T. Odate

1. Regional variations of zooplankton and phytoplankton with special reference to size distribution

Net plankton samples were collected at 14 stations by a NORPAC twin net (XX13 and GG54) tow from 200 m to the surface. Water samples were taken with Van Dorn samplers from several depths between 0 and 200 m. Phytoplankton in the water samples were fixed with Lugol's solution. Microzooplankton were concentrated by filtering 10 or 20 liters of seawater samples through a NXXX25 screen (40 μm mesh). Species and cell numbers of phytoplankters and microzooplankters will be determined later. Nutrient concentrations in ambient seawater were determined on board the ship. An aliquot of seawater was filtered through a Nuclepore filter (10 or 2 μm) or a Whatman GF/F filter, and chlorophyll a and phaeopigment were determined.

During sailing, 102 surface seawater samples were collected at about four-hour intervals from 4 m depth. The samples obtained were treated in the same way as described above. [T. Odate]

2. Studies on the behavior of organic matter in seawater and the processes of production and decomposition of organic substances

Seawater samples were collected from the surface to 5000 m at Stations B, C, D and F with Niskin bottles. The samples were filtered through a Whatman GF/F filter. The filtrates, to which 10^{-4}M of HgCl_2 was added were saved for later determinations of dissolved free amino acids and dissolved combined amino acids. Particulate samples collected on the filters were stored frozen at -20°C. Concentrations of ATP in the upper 200 m layer at Stations 1, 2, 3 and 5 of the Bering Sea were determined by the method described by Parsons et al. (1984) using 250 ml of water samples collected with Van Dorn bottles. Experiments were conducted at Station F to obtain information on the rate of zooplankton decomposition. Zooplankton samples were freeze-dried immediately after sampling, placed in a glass bottle containing 5 l of surface seawater, and incubated in the dark in a deck tank containing running surface

seawater. After 1.5, 5.5, 15, 30, and 50 hours, aliquots were taken out and filtered through a whatman GF/F filter. The amount and composition of amino acids in the filtrate will be examined later.

The analyses will be carried out in the shore laboratory after the cruise. [K. Tada]

A part of this research program was planned by Prof. Y. Maita, Faculty of Fisheries, Hokkaido University.

Reference

Parsons, T. R., Y. Maita and C. M. Lalli (1984). A manual of chemical and biological methods for seawater analysis. pp. 173, Pergamon Press, Oxford.

Fluorocytometry of oceanic particulate matter

T. Saino, K. Furuya and A. Hattori

Oceanic particulate materials were analysed with a JASCO FCS-1 cell sorter. For each particle in seawater (<40 μm), a 488 nm Ar-ion laser beam (500 mW) was irradiated, and the forward scattered light, right angle scattered light, and green and red fluorescence were monitored simultaneously.

Two fluorescent spherical latex beads (2 and 5 μm diameter) were used as standards for test of machine stability and as measure of each signal. A part of each water sample was saved for later image analysis by fluorescence microscopy (K. Furuya).

Data were obtained from shallow depths (<200 m) at Stations C, 1, 2, 3, 4, 5, 6, D, 7, 8, E, F, G, C' and H.

Estimate of cellular carbon in natural populations of phytoplankton

K. Furuya and T. Saino

The relationship between cellular carbon content and cell volume of natural populations of phytoplankton was investigated. Phytoplankton cells from the surface seawater were sorted using a JASKO FCS-1 cell sorter. Forward scattered light and red fluorescence were selected as indices. The sorted phytoplankton samples were kept frozen for later analysis of the carbon content with an CHN analyzer coupled to a quadrupole mass spectrometer. Care was taken to reduce a carbon blank.

Aliquots of the sorted samples were fixed with glutaraldehyde and processed following the procedure of Tsuji and Yanagita (1981). The cell volume of the phytoplankton and possible contamination of the other detrital particles were examined by epifluorescence microscopy.

Reference

- Tsuji, T. and T. Yanagita (1981). Improved fluorescence microscopy for measuring the standing stock of phytoplankton including fragile components. *Mar. Biol.*, 64, 207-211.

Composition and biomass of phytoplankton

K. Furuya

Size distribution of phytoplankton was investigated following the method of Tsuji and Yanagita (1981). Water samples collected from various depths shallower than 200 m were fixed with 1 % glutaraldehyde, and filtered through Gelman GA6 membrane filters. The phytoplankton collected on the filters were examined by fluorescence microscopy. Cell numbers and size distribution were determined with an image analyzer system fitted with an epifluorescence microscope (excitation: 420-490 nm). Using the data obtained, total cell volumes of various size groups of phytoplankton were calculated (Furuya 1982). Numbers of phycoerythrin-containing cells were separately counted by exciting the phytoplankton specimens with a green light of 520-550 nm. Biomasses of the individual size groups of phytoplankton in terms of C were estimated from the cell volumes using reported values for the volume-carbon conversion factor. Classification at a class level was also conducted based on observations under an ordinary microscope.

Phytoplankton species will be identified based on the information obtained by observation under an inverted microscope and by scanning electron microscopy.

References

- Furuya, K. (1982). Measurement of phytoplankton standing stock using an image analyzer system. Bull. Plankton Soc. Japan, 29, 131-132.
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Vertical distribution of submicron particles

I. Koike

Vertical distributions of submicron particles in the Bering Sea and northern North Pacific Ocean were determined. Discrete water samples were collected with a rosette sampler (5 l) or with 23-l Niskin bottles. At Stations C and F, bottom waters were also collected with the aid of a Pinger system.

An Elzone particle counter (80XY, Particle Data, Inc. Illinois, U.S.A.) fitted with a 12 μm orifice tube and a 10 μl volumetric tube was used. Nucleopore (0.2 μm) filtered 3.5 % NaCl solution served as a control. When necessary, the sample seawater was diluted with the same NaCl solution. Using 120 channels of the machine, particle counting can be made over the size range from 0.34 to 7.85 μm . However, we adopted values only in the size range from 0.38 to 2.02 μm (60 channels) because of high blank values in the range smaller than 0.38 μm and large fluctuation of signals in the range larger than 2.0 μm . The measurements were repeated at least three times and averaged to express numbers of particles in each size fraction. Total volume of the particles in each size fraction was calculated by using the particle numbers and assumed particle diameter.

Growth and carbon conversion efficiency of bacterial populations
in the euphotic layer of the northern North Pacific

I. Koike and L. Tupas

Bacterial growth in seawater was estimated from increase in particle number or volume in given size fractions during incubation. Sample seawater collected from the euphotic layer of the northern North Pacific was filtered through a Whatman GF/F filter (pore size ca. 0.6 μm) to eliminate the bacterial grazers. The filtrate (250 ml) was placed in a 500 ml glass bottle and incubated at in situ temperature in the dark. At 10-30 hr intervals, 30 ml portions were taken out, and introduced in a test tube (10 ml, for particle counting) and a plastic bottle (20 ml, for nutrient analysis).

The subsamples for particle counting were treated with Gelman disk (0.45 μm) filtered formalin (2 % final concentration). The change in the size distribution of the particles was monitored with an Elzone particle counter for 2 weeks. The subsamples for nutrients and other organic nitrogen and carbon measurements were stored frozen for later analyses.

Excreta from zooplankton were obtained by incubating copepods or sagitta in GF/F filtered surface seawater for 20 hrs. After the incubation, zooplankton were removed by gentle filtration through a 100 μm net. The filtrate was further incubated as described above.

About 5 l of GF/F filtered seawater with or without added zooplankton was prepared. The water sample was transferred, by using a siphon, into a 500 ml glass bottle. After allowing ca. 500 ml to overflow, the bottle was capped carefully and incubated at in situ temperature in darkness. Four glass bottles were prepared for one experiment. At 10-30 hour intervals, the bottles were sacrificed, one by one, for the determinations of oxygen, particle numbers and nutrients. The Winkler titration method using a 100 ml oxygen bottle was used for oxygen analysis. Carbon conversion efficiency during bacterial growth was calculated from the increase in particle volume and oxygen consumption by assuming appropriate conversion factors for cell carbon/volume and respiratory quotient.

Excretion of organic and inorganic compounds by
macrozooplankton and the utilization of excreta
by heterotrophic marine bacteria

L. Tupas and I. Koike

Experiments were performed on board to determine the dissolved organic and inorganic compounds excreted by the different macrozooplankton collected from stations in the northern North Pacific Ocean and the Bering Sea and also to study the utilization of these excreted dissolved compounds by heterotrophic marine bacteria.

Live zooplankton were collected with a 0.493 μm mesh size net with a mouth diameter of 0.5 m and fitted with a large cod end (2 liters capacity). Samples were collected from a depth of 200 m and the animals placed in a large container for sorting. Animals used for experiments were from the Amphipoda, Euphausiacea, Copepoda, Chaetognatha, Ctenophora, Schyphomedusae, and Thaliacea. Live animals were picked up one by one by gentle suction using an automatic pipet with a modified tip and transferred to 1 liter polycarbonate bottles containing 500 ml of either filtered sea water or synthetic sea water (0.45 μm Millipore HA). Incubation bottles contained from 5 to 50 animals depending on the size of the animals being incubated (the larger the animal, the less number incubated). The bottles were covered with aluminum foil and incubated in the dark at in situ temperature (4–6°C) for 24 hours. Forty ml of water was collected at 6-hour intervals and frozen for chemical analysis. Samples were collected with a siphon covered with a 100 μm net at the end.

To determine the utilization of excreta, animals were collected and treated as above and kept in the incubation bottles. After 24 hours the incubation water was transferred into another clean bottle using a siphon covered with a 100 μm mesh size net at the end to prevent the animals from being transferred into the new bottle and incubated further for 6 days under the conditions mentioned above. Forty ml of water (10 ml for microbial and particle counting; 30 ml for chemical analysis) was collected at 2-day intervals.

Feeding behavior of *Calanus plumchrus* and *Calanus cristatus*

H. Ishii and S. Nishizawa

The gut fluorescence method of Mackas and Bohrer (1976) is becoming popular in estimating the ingestion rate of herbivorous zooplankton under field conditions (Dagg and Grill, 1980). Gut clearance time is needed in calculation of the ingestion rate, but the effects of the particle concentration and feeding prehistory on the gut clearance time have not been considered in most previous studies. We tried to estimate the ingestion rates of freshly caught copepods, *Calanus plumchrus* and *C. cristatus*, which are the dominant species in the Bering Sea and the Oyashio Water, using the gut fluorescence method. We also aimed to clarify the effects of feeding prehistory on ingestion rate, the relation between diel vertical migration and feeding behavior, and the daily energy requirement of these copepods.

1. Zooplankton sampling

Zooplankton were collected from three depth zones, 0-50, 50-100, and 100-200 m, by divided vertical tows of a closing net (56 cm diameter, 493 μm mesh) with a 2-l cod-end. A portion of the samples was preserved for later analysis, and the remainder was used for immediate sorting of zooplankton. Copepodite V specimens of *C. plumchrus* and *C. cristatus* were picked up. These were ground in a 90 % acetone solution, and chlorophylls and phaeopigments were measured with a Turner Designs fluorometer. The residuals were frozen for later elemental analysis.

2. Feeding experiment

The zooplankton collected were reared in seawater at surface seawater temperature and acclimated. Starved specimens were prepared by leaving the zooplankton in GF/C filtered seawater for 1.5 days. The zooplankton samples with different feeding prehistory were prepared by rearing zooplankton in seawater with various particle concentrations for 1 day. The starved zooplankton (2 to 4 individuals for *C. plumchrus* and 1 for *C. cristatus*) were transferred to 100 ml glass bottles containing seawater with different particle concentrations. The other groups were transferred to 100 ml glass bottles filled with natural seawater containing the same particle concentrations as those used for the acclima-

tion. The experimental bottles were placed on a wheel rotator and incubated for 3 hours at 4.5°C in the dark. Initial and final particle concentrations were determined with an Elzone electronic particle counter of Particle Data Inc. using a 120 µm orifice. The ingestion and filtering rates were calculated using the equation of Frost (1972). After the experiments, the gut fluorescence was measured, and the gut clearance time was calculated. The rates of respiration and ammonia excretion were estimated from the differences in oxygen and ammonia concentrations before and after 24 hours incubation.

References

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Ecological study of zooplankton and micronekton

M. Terazaki and H. L. Suh

1. Collection of zooplankton and micronekton

A NORPAC-twin net consisting of 0.10 mm and 0.33 mm-mesh nets was towed vertically from a depth of 150 m to the surface at each station. The tows were carried out mostly at night to collect epipelagic zooplankton.

A Motoda horizontal net (MTD net) was towed at 3-4 hour intervals in 5 different strata between 200 m and the surface at Station C. The net tows were also made at Stations 1 (0-200 m), 2 (0-80 m) and 3 (0-80 m).

Large zooplankton and micronekton were collected by an oblique tow of an ORI net (0.69 mm-mesh) in the layer between 1470 m and the surface at Stations B, C and F.

An ORI type Multiple Plankton Sampler (ORI-MPS) with 0.33 mm-mesh nets was towed vertically at 6-hour intervals in 4 different strata between 1500 m and the surface at Station B. Large zooplankton, especially chaetognaths and copepods were collected. The ORI-MPS was also towed, day and night, in 3 different strata between 2000 m and the surface at Station F'.

2. Study of diurnal migration of zooplankton

Diurnal migration of amphipods (*Parathemisto japonica* and *P. pacifica*), euphausiids (*Euphausia pacifica* and *Thysanoessa longipes*) and chaetognaths (*Sagitta elegans* and *Eukrhonia hamata*) was investigated using the specimens collected with MTD nets. Spatial and temporal variations of the distribution of meso- and bathypelagic zooplankton were examined using the samples collected with the ORI-MPS.

3. Feeding habits of amphipods and euphausiids

The food items and feeding rhythm of amphipods (*P. japonica* and *P. pacifica*) and euphausiids (*T. longipes*, *T. inermis*, *T. raschi*, *T. spinifera* and *E. pacifica*) were analyzed using the samples collected with MTD nets. Pigment analysis of stomach contents was also conducted.