# NOAA Data Report ERL PMEL-37

#### PUGET SOUND SEDIMENT TRAP DATA: 1980-1985

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# CONTENTS

1.0 INTRODUCTION
2.0 METHODS 4
2.1 Collection and Processing 4
2.2 Analyses
2.2.1 Total Sedimentation Rate 4
2.2.2 Size Fractionation
2.2.3 Pigments
2.2.4 Organic Carbon and Nitrogen 7
2.2.5 Trace Metals 7
2.2.5.1 Trace Metals – 1981 7
2.2.5.2 Trace Metals – 1982 8
2.2.5.3 Trace Metals – 1983 to 1984
2.2.5.4 Trace Metals – 1985 8
2.2.6 Trace Organics 8
3.0 RESULTS 12
3.1 1981 Sampling Year 12
3.2 1982 Sampling Year
3.3 1983 Sampling Year
3.4 1984 Sampling Year
3.5 1985 Sampling Year
4.0 ACKNOWLEDGMENTS
5.0 REFERENCES
6.0 PUGET SOUND BIBLIOGRAPHY 29
APPENDIX (on microfiche in pocket inside back cover)

# **FIGURES**

1.	Mechanical design of the sediment trap	. 5
2.	Fractionation scheme for sediment trap particles	. 6
	TABLES	
1.	Summary of sediment traps recovered	. 3
2.	Quality control/quality assurance data for trace metals	. 9
3.	Organic recovery data (%); 1981	11
4.	Sediment trap location and sampling data for 1981	13
5.	Availability of sediment trap data for 1981	14
6.	Sediment trap location and sampling data for 1982	16
7.	Availability of sediment trap data for 1982	17
8.	Sediment trap location and sampling data for 1983	19
9.	Availability of sediment trap data for 1983	20
10.	Sediment trap location and sampling data for 1984	22
11.	Availability of sediment trap data for 1984	23
12.	Sediment trap location and sampling data for 1985	25
13.	Availability of sediment tran data for 1985	25

# Puget Sound Sediment Trap Data: 1980-1985

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#### 1.0 INTRODUCTION

In 1979, scientists at the Pacific Marine Environmental Laboratory began investigating the sources, transformation, transport and fate of pollutants in Puget Sound and its watershed under Sec. 202 of the Marine Protection, Research and Sanctuaries Act of 1971 (P.L. 92-532) which called in part for "...a comprehensive and continuing program of research with respect to the possible long range effects of pollution, overfishing, and man-induced changes of ocean ecosystems..." The effort was called the Long-Range Effects Research Program (L-RERP) after language in the Act and was later called the PMEL Marine Environmental Quality Program. Building on research then underway at PMEL on estuarine circulation, laboratory scientists began a coordinated study that began with the description of the distribution of properties (salinity, temperature, trace metals and trace organics) in the water column and underlying sediments. The objectives of the Marine Environmental Quality Trace Metal and Organics Program were 1) to quantify the sources and sinks of selected trace metals and organics for Puget Sound, 2) to determine geochemical mechanisms that transform trace metals between the dissolved and particulate phases and 3) to determine to what extent these geochemical mechanisms alter the fate of trace metals and organics entering Puget Sound. Work began in rivers discharging into Puget Sound and process studies were undertaken to understand the role of flocculation in trace metal transport. Subsequently the research centered on the role of suspended sediments in transporting and redistributing trace metals and organics in the main basin of the Sound. Research activities included deployment of long-term current meter moorings, acquisition of a library of sediment cores, deployment of sediment traps and the analyses of dissolved and particulate chemical constituents of the water column and sediments. The scientific results of these activities have been reported in over 100 publications (see Section 6).

The Long-Range Effect Research Program consisted of 1) sampling dissolved and particulate constituents in the water column by bottle sampling, 2) sampling settling particles by sediment trap and 3) sampling sediments by grab, box, gravity and Kasten corers. All three type of samples collected between 1979 and 1985 for the two urban embayments (Elliott and Commencement Bays) are reported in Paulson et al. (1991a). All the water column data in the

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open waters of Puget Sound collected between 1981 and 1985 are reported in Paulson et al. (1991 b). Water column data collected from a single cruise in 1986 in both the open waters and embayments of Puget Sound are reported in Paulson et al. (1991c). In this Data Report, a variety of data from particles collected in 104 traps deployed on 34 moorings in open waters between 1980 and 1985 are presented (Table 1).

Because these measurements constitute the most extensive data base of trace metal and trace organic observations in Puget Sound, many of which have been unavailable to other investigators, we feel that they should be widely available to the local scientific community as well as others interested in estuarine geochemistry. The text of this data report begins with the sampling and analytical methods with the accompanying quality control/quality assurance data. The text of the data sections are a summary of the available data and published literature in which the data is interpreted along with a catalogue of the data available in the Appendix (on microfiche located in the back pocket of this data report).

A catalogue of sediment data for the open waters of Puget Sound is not available at this time. Therefore, below you will find a description of citations in which PMEL's sediments collected by PMEL's L-RERP sampling program have been analyzed and reported. Sedimentation rates in Puget Sound are reported in Lavelle et al. (1985, 1986). Baker et al. (1985) discuss the carbon and pigment content and sedimentation rates in Puget Sound. Trace metal content, geochemistry and sedimentation rates are discussed in Romberg et al. (1984), Feely et al. (1986), Bloom and Crecelius (1987) and Paulson et al. (1988a). Bates et al. (1984, 1987) and Murphy et al. (1988) present and discuss trace organic content in the sediments. All ancillary data collected from the cores associated with the publications cited above have been submitted in uncollated form to the data management team of the Puget Sound Water Quality Authority.

Table 1. Summary of Sediment Traps Recovered

Region	Time series in CMB	AI, CMB, EP	Transect	ACTOSS EF, COL EP AI, CMB, EP AI, CMP, EP	SJF, AI, CMB,	AI, CMB, EP	CMB
Dates (Inclusive)	5 Dec 80 – 2 Feb 81 2 Feb 81 – 16 Apr 81 22 Apr 81 – 5 Jul 81 21 Jul 81 – 29 Sep 81 7 Oct 81 – 19 Dec 81	Feb 82 – 25 Apr 82	18 Mar 82 – 24 Apr 82	13 Apr 83 – 29 Apr 83 26 Apr 83 – 29 Aug 83 29 Aug 83 – 7 Dec 83	1 Dec 83 – 30 May 84	5 Apr 84 – 23 Jul 84	29 Mar 85 – 22 Jun 85 10 Apr 85 – 12 Apr 85
Cylinder Collection Time (hr)	144–168	137–168	91	24 288 228	288	264	204
Number of Traps	22	17	21	०००	16	9	1 3
Number of Moorings	S	v	7	H & &	9	ю	1
Sampling Year	1981	1982	1983		1984		1985

SJF Strait of Juan de Fuca
Al Admiralty Inlet
CMB Central Main Basin
EP East Passage
WB Whidbey Basin
COL Colvos Passage

# 2.0 METHODS

#### 2.1 Collection and Processing

Settling particles were trapped using the cylindrical Sequentially Sampling Sediment Trap described by Baker and Milburn (1983). The smooth polyethylene collection funnel has a collection cross-sectional area of 324.3 cm<sup>2</sup> and an aspect ratio of 3.0:1 (Fig. 1). Settling particles collect in the funnel and are directed in turn to one of 10 acrylic containers (200 ml each). The acrylic containers have been filled with a dense filling solution to minimize loss of particles by turbulence. Toxins such as sodium azide are added to the dense filling solution brine to prevent organic growth. After a preset time, the next acrylic container on the sampling unit is rotated under the funnel. The sediment traps are kept upright during collection and are closed during the recovery process. In each sampling year, sediment traps were deployed in different locations throughout Puget Sound for different durations. A summary of the number and duration of deployments for each sampling year is presented in Table 1.

Upon recovery, the sampling containers were transported to the laboratory and the contents split. Approximately half of the supernatant (100 ml) was first decanted from the sampling containers. This liquid was stored in high-density linear polyethylene bottles or discarded. The remaining sample of particles and supernatant was then thoroughly mixed and split into separate aliquots for measurement of total dry weight, pigments, total carbon and nitrogen, trace metals and trace organic measurements.

#### 2.2 Analyses

#### 2.2.1 Total Sedimentation Rate

The contents of each aliquot for total dry weight analysis were filtered onto a pre-tared, 47 mm, 0.2  $\mu$ m Nuclepore filter and vacuum desiccated. After at least a week of drying, the filters were re-weighed on electrobalances. The total dry weight was calculated by difference. Sedimentation rates were calculated from the total dry weight, the duration of the collection period and the collection cross-sectional area.

#### 2.2.2 Size Fractionation

Particles were separated according to size and density and collected on 0.4  $\mu$ m membrane filters for weighing. A separation scheme (Fig. 2) was designed to isolate fine and coarse mineral grains, fecal pellets and coarse low-density particles. In step (1), aliquots from each trap were gently washed through a 38  $\mu$ m sieve. The large particles retained on the sieve were then segregated at step (2) into density classes by flotation in carbon tetrachloride with density  $\rho = 1.6 \text{ g/cm}^3$  (Dillon, 1964). The low-density fraction at step (3) was vigorously rewashed through a 38  $\mu$ m sieve using hot soapy water to disaggregate fecal pellets. Sedimented particles were density segregated by centrifugation on a saturated zinc chloride solution ( $\rho = 1.9 \text{ g/cm}^3$ ).

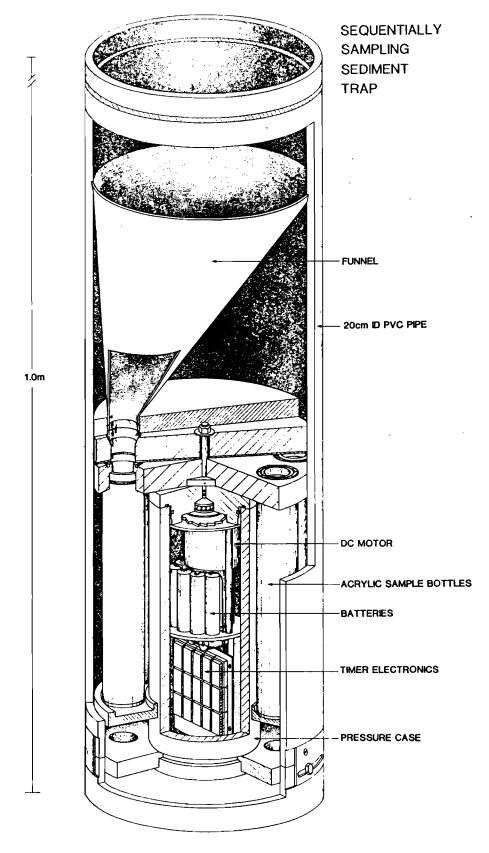


Fig. 1. Mechanical design of the sediment trap. The electronic timing components are accessible after removing the pressure-case end cap at the base of the unit.

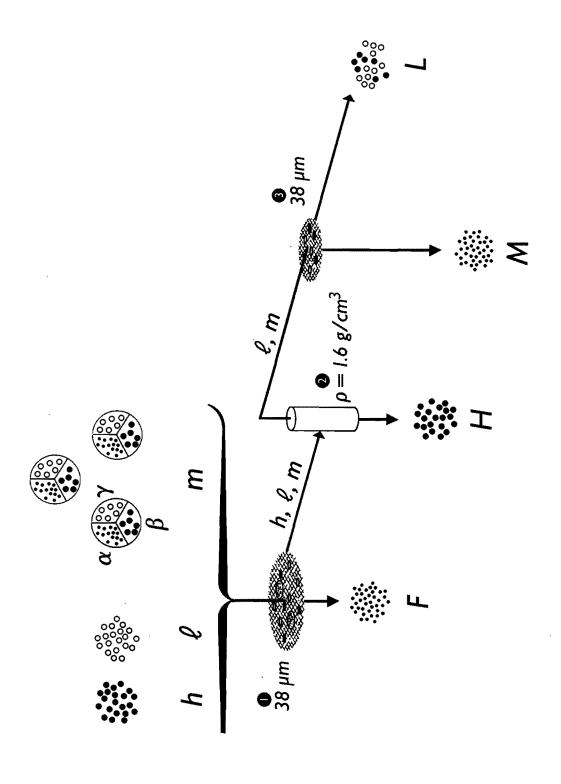


Fig. 2. Fractionation scheme for sediment trap particles. At the first step, fine particles F were mechanically separated by washing through a 38 µm sieve. density coarse particles L and intact macroaggregates were vigorously rewashed through a 38  $\mu$ m sieve. The fine fraction ( $\alpha$ ) of the disaggregated macroaggregates passing through the sieve was collected as fraction M. The particles retained on the sieve at step (3) consisted of low-density coarse Particles retained on the sieve were segregated according to density at step (2), with high-density coarse particles H being separated out. At step (3), lowparticles, including the fraction y disaggregated from macroaggregates, as well as the high-density coarse fraction  $\beta$  disaggregated from macroaggregates.

#### 2.2.3 Pigments

Pigments in trapped particles were measured in aliquots filtered onto glass-fiber filters, stored frozen, and analyzed by fluorometry as described by Holm-Hansen *et al.* (1965). The total pigment in a sample (chl<sub>T</sub>) is expressed in terms of chlorophyll a equivalents, calculated as chlorophyll a + (1.51) pheophorbide (Shuman and Lorenzen, 1975).

### 2.2.4 Organic Carbon and Nitrogen

Total particulate C and N was measured in aliquots filtered onto silver filters, stored frozen, and combusted in a Perkin-Elmer 240B CHN analyzer. Acid treatment of selected samples indicates that particulate C was typically 95% organic.

#### 2.2.5 Trace Metals

Several different analytical procedures were used to determine the trace metal content of the settling particles for each sampling period.

2.2.5.1 Trace Metals – 1981. The trace metal aliquots from the ten cylinders from each sediment trap were combined to give one sample representing the entire trap deployment. The combined sample was then sieved through a 64  $\mu$ m polyethylene screen and the <64  $\mu$ m material was collected on a 142 or 47 mm, 0.4  $\mu$ m Nuclepore filter and vacuum desiccated. The <64  $\mu$ m fraction was analyzed for trace metals.

Trace elements associated with the poorly structured hydrous oxide phases of the suspended matter and sediment trap samples were determined by the method of Chester and Hughes (1967) with slight modifications. Desiccated samples were leached with 5 ml of a mixed reagent containing 0.04 hydroxylamine hydrochloride in 25% Ultrex® acetic acid at 50°C for 2 h. The resulting supernatant was filtered through an acid-cleaned polypropylene-glass filtration unit containing a 0.2  $\mu$ m Nucleopore filter. The residue was rinsed with quartz-distilled water, then filtered; the supernatant was then combined with the original supernatant, acidified with 0.5 ml of concentrated Ultrex® HCl diluted to 20 g weight, and analyzed for trace elements by the graphite furnace atomic absorption spectrometer (GFAAS) procedures described by Feely et al. (1981).

Trace elements associated with particulate organic matter were determined using the procedures of Landing and Feely (1981). Separate 100-mg aliquots of sediment trap material were heated in 10%  $H_2O_2$  for 18 h at 65°C and then sonicated with continued heating for another 24 h. The samples were decanted and filtered through an acid-cleaned 0.2- $\mu$ m Nucleopore filter. The samples were rinsed with two portions of quartz-distilled water, acidified with 0.5 ml of concentrated Ultrex® HCl, diluted with quartz-distilled water to 20 g total weight, and analyzed for trace elements by flameless atomic absorption.

Total metal concentrations for Al, Fe, Mn, Ni, Cu, Zn, Pb, Cr, Cd and Co in OMPA1 and OMPA4 samples were determined by dissolution in a HF-HCl-HNO<sub>3</sub> acid digestion according to the method of Eggiman and Betzer (1976) followed by analysis of the digested acid solution by GFAAS. The precision of total analysis of samples is given in Table 2. Arsenic in OMPA1 samples and Fe, Mn, Cu, Zn, Pb, Cr, V and As in OMPA2 and OMPA3 samples were analyzed by XRF according to the method of Nielson (1977). Ag, Se and Hg were determined according to the method of Bloom and Crecelius (1987).

- 2.2.5.2 Trace Metals 1982. The trace metal aliquots from the ten cylinders from each trap were combined in the manner described for the 1981 trace metal samples. Total Cu, Pb, Ag and Hg were determined by the method of Bloom and Crecelius (1984).
- 2.2.5.3 Trace Metals 1983 to 1984. The trace metal aliquots from the ten cylinders from each trap were combined and processed in the manner described for the 1981 trace metal samples. Total Mn, Cu, Pb and Cd concentrations were determined by dissolution in a HF-HCl-HNO<sub>3</sub> acid digestion according to the method of Eggiman and Betzer (1976) followed by analyses of the digested acid solution by GFAAS. Quality assurance/quality control data for this set of analyses is given in Table 2.
- 2.2.5.4 Trace Metals 1985. For the 1985 sediment trap deployments, trace metal aliquots from each cylinder were filtered onto separate filters. The Mn concentrations in <63  $\mu$ m fraction and the >63  $\mu$ m fraction for each cylinder were analyzed separately. The Mn associated with the oxide phase of the sediment trap particles was selectively extracted with hydroxylamine hydrochloride according to the method of Robbins et al. (1984). The extractant from the selective extraction procedure was analyzed by GFAAS. The Mn in the residual phase was determined by the dissolution in a HF-HCl-HNO<sub>3</sub> acid digestion according to the method of Eggiman and Betzer (1976) followed by analyses of the digested acid solution by GFAAS.

# 2.2.6 Trace Organics

In most cases, the trace organic aliquots from each cylinder were combined and processed in the manner described for the 1981 trace metal samples. In some cases, a sediment trap contained enough material to combine aliquots from less than the ten cylinders (usually 3–4 cylinders) to produce several samples from the same sediment trap.

The analytical procedure for trace organics included Soxhlet extraction with methanol/dichloromethane, silica gel adsorption chromatography to separate the aliphatic and aromatic fractions, and Sephadex gel permeation chromatography to further concentrate the polycyclic aromatic hydrocarbons (PAH) (Hamilton et al., 1984). Compounds were quantified with a Hewlett-Packard 5880 gas chromatograph and a Hewlett-Packard 5993 gas chromatograph-

Table 2. Quality Control/Quality Assurance Data for Trace Metals

රි	5–10%	19 16	36 36	25					
ප					0.15	19	0.35 ±0.07 0.25 ±0.07	0.33 ±0.06 N/A	0.60 ±0.01 0.59 ±0.1
Pb	6–16%	33	15 14	21 27	7	1	2121	844	73. 14. 14. 14.
Zn	3-25%	51 86	110 125	153 140					
ರೆ	4%	88	22 16	33	7	6	18 13 13	28 27	£ 25 £ 3
ï	10-11%	7	19 10	74					
Mn	0.5-4%	942 774	1660 1400	935 774	12	4	227 ±13 229 ±15	651 ±15 770	472 ±78 513 ±25
Fe	<sup>7</sup> 4%								
¥	1–6%	6.9	5.4	7.1					
	Sample Precision (%CV; n=3)	AGV SRM Observed AGV SRM Literature	BCR SRM Observed BCR SRM Literature	MAG SRM Observed MAG SRM Literature	Determination Limit (ppm)	Sample Precision (%CV, n=3)	BCSS-1 SRM Observed BCSS-1 SRM Certified	MAG SRM Observed MAG SRM Literature	MESS-1 SRM Observed MESS-1 SRM Certified
	1981				1982–1984				

mass spectrometer using either an OV-1 (aliphatic) or SE-54 (PAH) fused silica column with temperature programmed from 70 to 270°C at 4°C/min. Recovery standards were used to correct volatilization losses (Barrick et al., 1980) (Table 3). Analytical precision for the individual hydrocarbons was  $\pm$  10% (one standard deviation expressed as a percentage of the mean) and the limit of detection for each individual compound was generally less than 5  $\mu$ g/g. PAH compounds derived from combustion processes are designated combustion PAH (COMB PAH) and represent the sum of nine compounds—fluoranthene, pyrene, benz[a]anthracene, chrysene, benzofluoranthenes, benzo[e]pyrene, benzo[a]pyrene, indeno[c,d]pyrene, and benzoperylene. All concentrations are expressed in nanograms per gram dry weight of sediment.

Table 3. Organic Recovery Data (%); 1981

	Alphatic	<u>PAH</u>
OMPA 1 range	73 ± 4 (68–78)	54 ± 6 (48–56)
OMPA 2 range	97 ± 7 (85 ± 100)	55 ± 12 (42–67)
OMPA 4 range	85 ± 14 (67–99)	93 ± 33 (78–139)
OMPA 5 range	10 ± 6 (6–14)	83 ± 19 (40–103)

#### 3.0 RESULTS

The data from recovered sediment traps is organized by sampling year in the following sections. For each section, a brief description of the locations and deployment intervals of the sediment traps for the sampling year is given. This is followed by the citations where interpretations of the data can be found. Each section also contains two tables. The first table gives sampling data for each sediment trap in the following order: sediment trap name, serial number (S/N), mooring name, latitude, longitude, water depth, depth of sediment trap deployment, collection start time, duration of collection time for each cylinder, collection stop time and comments. The second table indicates what type of data is available in the Appendix for each sediment trap such as pigment, carbon and nitrogen content, size fractionation, trace metal (TM) or trace organic (Org.) data. The last column of this table indicates the page number in the Appendix that a particular type of data for the given trap can be found. The Appendix is organized in sets. Firstly, individual tables for each sediment trap within the set are presented with data for the ten cylinders organized by rows. The total sedimentation rate is always given for each cylinder. Pigment, carbon and nitrogen and size fractionation data for each cyclinder can also found in these tables for the individual sediment traps, if available. For trace metal and trace organic analysis, the samples from the ten cylinders from each trap were usually combined. Therefore, the data for trace metals and trace organics from each set of sediment traps is presented in two tables in the Appendix following the set of individual sediment trap tables. In the table for trace metal and trace organic data, data from each sediment trap is given by rows.

#### 3.1 1981 Sampling Year

The 1981 sampling program consisted of a time series at a central main basin location with almost continuous sampling between 5 December 1980 and 19 December 1981 (Table 4). Usually traps were deployed at depths of 50, 100, 160 and 200 m. During the summer deployment (OMPA4), a trap was placed at 20 m.

The total sedimentation rates, pigment and carbon data (Table 5) are presented and discussed in Baker et al. (1985). Feely et al. (1986) presents and discusses the data for total trace metal content and for trace metal content of electively extracted phases of the settling particles. Further discussion of the trace metal data can be found in Paulson et al. (1988a). Mn concentrations on settling particles at the central main basin location are discussed in Cudaback et al. (1991). The trace organic data is presented and discussed in Bates et al. (1984, 1987) and Murphy et al. (1988).

Table 4. Sediment Trap Location and Sampling Data for 1981

10 Comments	1 of Triple		1 of Triple	Averaged	Averaged
	81 00 81 00 81 00	81 00:00 81 02:00 81 02:00	81 02 81 12 81 12 81 12 12 12	81 02:00 81 02:00 81 02:00 81 02:00	DEC 81 00:00
9 Stop: Date/	000				0 0 0 0 0
8 Duration Cylinder	144	11 16 16 16 16 16 16 16 16 16 16 16 16 1	161 168 168 168	161 161 161 161	176 176 176 176 176
t: /	888	8188	8 8 8 8 1 8 1 8	81 81 81 81	81 16:00 81 16:00 81 16:00 81 16:00 81 16:00
7 Start Date/	5 DEC 5 DEC 5 DEC		2 FEB 22 APR 22 APR 22 APR 22 APR		7 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
6 Sample Depth	50 100 160	205 42 42 165	197 50 100 160 200	20 100 160 200	20 50 100 160 200
5 Water Depth	210 210 210	202 202 203 203 203 203	550 0 550 550 550 550 550 550 550 550 5	210 210 210 210	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
4 Long.					122 27.3 122 27.3 122 27.3 122 27.3 122 27.3
3 Lat.	47 41.9 47 41.9 47 41.9	47 41.9	47 41.5 47 41.5 47 41.5 41.5 41.5	47 41.7 47 41.7 47 41.7 47 41.7	47 41.7 47 41.7 47 41.7 47 41.7
2 Mooring 3 Lat.	OMPA1 OMPA1 OMPA1	OMPA1	OMPA2 OMPA3 OMPA3 OMPA3	OMBA4 OMBA4 OMBA4 OMBA4	OMPAS OMPAS OMPAS OMPAS
1 S/N	491	10 H 01	るこのの4	3£7 5 1	15.4 3 15.2 7.5 4 3 15.2
O Trap Name	PS811-50 PS811-100 PS811-160	PS811-200 PS812-50 PS812-100	PS812-200 PS813-50 PS813-100 PS813-160 PS813-200	PS814-20 PS814-100 PS814-160 PS814-200	PS815-20 PS815-50 PS815-100 PS815-160 PS815-160

Table 5. Availability of Sediment Trap Data for 1981

0 Trap Name	1 S/N	2 Mooring	3 Pig- ments	4 Carbon	5 Size Frac.	6 TM	7 Org.	8 Page
PS811-50	4	OMPA1	x	x				6
PS811-50	-	VIII	_	_		X		10
PS811-50						_	x	11
PS811-100	6	OMPA1	x	x				7
PS811-100	•	VIII 111		-		x		10
PS811-100						_	x	11
PS811-160	1	OMPA1	x	x				-8
PS811-160	-	<b>VIII</b> 111		_		x		10
PS811-160							x	11
PS811-200	5	OMPA1	x ·	x				9
PS811-200	•	<b>Jan 111</b>		_		x		10
PS811-200						_	x	11
PS812-50	1	OMPA2	x	x	x			12
PS812-50	_		-				x	16
PS812-100	2	OMPA2	x	x	x		_	13
PS812-100	_	·				x		15
PS812-100							x	16
PS812-200	6	OMPA2	x	x	x		_	14
PS812-200	•	VIII 1111		-			x	16
PS813-50	1	OMPA3	x	x	x			17
PS813-50	-	VIII 110		_		x		21
PS813-50						-	x	22
PS813-100	2	OMPA3	x	· x	x			18
PS813-100	-	OLIL ILO	. 44	_			x	22
PS813-160	3	OMPA3	x	x	x		_	19
PS813-160	•	OLL PO		-		x		21
PS813-160						A	x	22
PS813-200	4	OMPA3	x	x	x			20
PS813-200	•	0111110		-	_	x		21
PS813-200						**	x	22
PS814-20	3&7	OMPA4	x	x	x			23
PS814-20				_		x		27
PS814-20							x	28
PS814-100	5	OMPA4	x	x	x			24
PS814-100	_					x		27
PS814-100						_	x	28
PS814-160	2	OMPA4	x	x	x		_	25
PS814-160	<del></del>				_	x		27
PS814-160							x	28
PS814-200	1	OMPA4	<b>x</b> .	X ·	x		_	26
PS814-200						x		27
PS814-200							x	28
PS815-20	1&2	OMPA5	x	x	x		-	29
PS815-20					<del>-</del>		x	34
PS815-50	3	OMPA5	x	x	x			30
PS815-50							x	34
PS815-100	4	OMPA5	x	x	x		_	31
PS815-100							x	34
PS815-160	5	OMPA5	x	x	x			32
PS815-160	•			_			X	34
PS815-200	7	OMPA5	x	x	x			33
PS815-200					_		x	34
								-

#### 3.2 1982 Sampling Year

The 1982 sampling program consisted of 5 moorings deployed between 8 February 1982 and 25 April 1982 (Table 6). In the Admiralty Inlet area, a mooring off Foulweather Bluff (PS821) held two sediment traps (50 and 100 m) while PS825 off Point No Point contained 4 sediment traps (19, 50, 90 and 175 m). A mooring at the central main basin location held two traps (175 and 195 m). In the northern end of East Passage, four sediment traps (20, 90, 147 and 177 m) were deployed on mooring PS8213. In Poverty Bay, mooring PS8214 held four sediment traps at approximately the same depth.

The trace organic data (Table 7) is presented and discussed in Bates et al. (1987) and Murphy et al. (1988).

Table 6. Sediment Trap Location and Sampling Data for 1982

10 Comments	of Triple
ī	15:00 115:00 115:00 115:00 115:00 02:00 02:00 02:00 02:00
	882211111111111111111111111111111111111
9 Stop: Date/	25 APR 25 APR
8 Duration Cylinder	168 168 168 168 168 1137 1137 1137 1137 1137
	17:00 13:00 13:00 13:00 13:00 15:00 15:00 14:00 14:00 14:00
7 Start: Date/	8 FEB 82 8 FEB 82 8 FEB 82 8 FEB 82 8 FEB 82 9 FEB 82 9 FEB 82 25 FEB 82 25 FEB 82 25 FEB 82 24 FEB 82 24 FEB 82 24 FEB 82
6 Sample Depth	100 100 50 119 175 1175 120 20 147 147 149
5 Water Depth	108 108 200 200 200 200 203 184 1184 1184 1175 1175
4 Long.	122 34.2 122 28.2 122 28.2 122 28.2 122 28.2 122 28.2 122 24.0 122 24.0 122 24.0 122 24.0 122 24.0 122 24.0 122 24.0
3 Lat.	44444444444444444444444444444444444444
1 S/N 2 Mooring 3 Lat.	PS821 PS821 PS825 PS825 PS825 PS825 PS825 PS8210 PS8213 PS8213 PS8213 PS8214 PS8214
1 S/N	2
O Trap Name	PS821-50 PS821-100 PS825-19 PS825-19 PS825-175 PS825-175 PS8210-175 PS8210-195 PS8213-20 PS8213-20 PS8213-147 PS8214-20 PS8214-10 PS8214-20 PS8214-10

Table 7. Availability of Sediment Trap Data for 1982

0 Trap Name	1 S/N	2 Mooring	3 Pig- ments	4 Carbon	5 Size Frac.	6 TM	7 Org.	8 Page
PS821-50 PS821-50	4	PS821	x			x		35 51
PS821-100	2	PS821	x			x		36 51
PS821-100 PS825-50	4	PS825				A	x	54
PS825-100 PS825-19	13	PS825	x				X	54 37
PS825-19	10	10020				x		51
PS825-19 PS825-90	6	PS825	x				x	54 38
PS825-90	ŭ	10020				x		51
PS825-90 PS825-175	12	PS825	x				x	54 39
PS825-175			<del></del>			x		51
PS825-175 PS825-195	7	PS825	x				х	54 40
PS825-195						x	••	51
PS825-195 PS8210-175	9	PS8210	x				x	54 41
PS8210-175 PS8210-195	11	D00010	x			x		51
PS8210-195 PS8210-195	11	PS8210				x		<b>42</b> 51
PS8213-20 PS8213-20	8	PS8213	X		X	x		43 51
PS8213-20							x	54
PS8213-90 PS8213-90	19	PS8213	x		X		x	44 54
PS8213-147	10	PS8213	x					45
PS8213-147 PS8213-147						х	x	51 5 <b>4</b>
PS8213-177	15	PS8213	x		x			46
PS8213-177 PS8213-177						x	x	51 54
PS8214-20	16	PS8214	x					47
PS8214-20 PS8214-20						X	x	51 5 <b>4</b>
PS8214-90	17	PS8214	x					48
PS8214-90 PS8214-90						x	x	51 54
PS8214-149	18	PS8214	x			v		49
PS8214-149 PS8214-149						X	x	51 5 <b>4</b>
PS8214-177 PS8214-177	20	PS8214	x			x		50 51
PS8214-177							x	54

# 3.3 1983 Sampling Year

The 1983 sampling program consisted of two sets of sediment trap deployments (Table 8). A short term experiment (18 March 1983 to 24 April 1983) emphasized sediment transport in East Passage. An across-axis transect consisting of 4 moorings (PS833–PS836) stretched across East Passage east of Three Tree Point. Each mooring held a sediment trap at 50 and 100 m while the center mooring (PS835) also contained sediment traps at 30, 148 and 214 m. Sediment traps on moorings in Poverty Bay (PS838; 50, 100 and 188 m), off Brown's Point (PS8310; 50, 100 and 177 m) and Colvos Passage (PS8312; 88 m) also were part of the short term experiment. A single mooring (PS8324) with a cylinder duration of one day was deployed in Poverty Bay with double sediment traps deployed at 30, 90 and 155.

Between 13 April 1983 and 29 August 1983, sediment trap were deployed at 30 and 100 m off Point No Point (PS8329), in central main basin (PS83-26) and off Three Tree (PS83-27). A second set of sediment traps at 30 and 100 m at the same locations were deployed between 29 August 1983 and 7 December 1983 (PS8333, PS8330 and PS8331, respectively).

Mn concentrations on settling particles at the central main basin location (Table 9) are discussed in Cudaback et al. (1991). Trace organic data for the 1983 sediment trap program is presented and discussed in Bates et al. (1987) and Murphy et al. (1988)

Table 8. Sediment Trap Location and Sampling Data for 1983

10 Comments	of Double	of					. of Triple	o£	of													Jo	of Double			of													
9 Stop: Date/	4 APR 83 22:00	4 APR	4 APR 83 22:00	4 APR 83	4 APR 83	4 APR	4 APR 83 22:00	4 APR 83	4 APR 83 22:00	24 APR 83 22:00	4 APR 83	4 APR 83	4 APR 83	4 APR 83	83	4 APR 83	4 APR 83	83	4 APR 83	4 APR 83	ARP 83	APR 83 00:00	APR 83 00:00	APR 83 00:00	APR 83 00:00	83	APR 83 00:00	AUG 83	AUG 83	29 AUG 83 00:00	AUG 83	AUG 83	9 AUG 83	7 DEC 83 00:00	DEC 83	83	DEC 8	ထ	m
8 Duration Cylinder	91	91	91	91	91	91	91	91	91	91	91	91	91	91	91	91	91	91	91	91	91	24	24	24	24	24	24	288	288	288	288	288	288	228	228	228	228	228	228
e 7 Start: Date/	18 MAR	18 MAR 83 00:	18 MAR 83	18 MAR 83	18 MAR	18 MAR	18 MAR 83	18 MAR	18 MAR 83	18 MAR 83	18 MAR 83	18 MAR 83	18 MAR 83	18 MAR	18 MAR	18 MAR 83	18 MAR	18 MAR 83	18 MAR 83	18 MAR	18 MAR 83	13 APR 83	13 APR	13 APR 83	13 APR 83	13 APR 83	13 APR 83	26 APR 83 1	26 APR 83	25 APR 83	25 APR 83	26 APR 83	26 APR 83	29 AUG 83 1	29 AUG 83 1	29 AUG 83 1	29 AUG 83 1	29 AUG	29 AUG 83 1
ar 6 Sample th Depth	 	77 50	-	33 50	-	221 3						221 214									174 88					173 15			202 10			198 30	•			33 30			98 100
5 Water Depth									3 22	3 221	3 221	m	7	7	_	_	_																						10
4 Long.	25	122 25.5	25	24	24	24	24	24	24	122 24.	-				122 22.(						122 31.2					122 22.0										22	122 22.4	22	22 28
3 Lat.	6 .	26.	26.	Ġ	26.	27.	27.	27.	27.	~	27.	27.	27.	27.	47 21.7	21.	21.	Н	47 19.7	-1	25.	21.	21.	٠	21.	7	ä	41,	41.	26.	26.	53	53.	41.	41.	ġ	4	53.	
2 Mooring	PS833	PS833	PS833	PS834	PS834	PS835	PS835	PS835	PS835	PS835	PS835	PS835	PS836	PS836	PS838	PS838	PS838	PS8310	PS8310	PS8310	PS8312	PS8324	PS8324	PS8324	PS8324	PS8324	8832	5832	5832	PS8327	\$832	5832	5832	8833	8833	8833	00	5833	5833
1 S/N	26		27	28	29	14	15	16	17	18	20	23	24	25	8	9	7	11	12	13	30	31	32	33	34	35	36	17	15	14	18	16	20	7	9	7	11	13	23
O Trap Name	33-	PS833-50B	PS833-100	PS834-50	PS834-100	PS835-31	PS835-50	PS835-50B	PS835-50C	PS835-100	PS835-148	PS835-214	PS836-50	PS836-100	PS838-50	PS838-100	PS838-166	PS8310-50	PS8310-100	PS8310-170	PS8312-88	PS8324-30	PS8324-30B	PS8324-90	PS8324-90B	S	PS8324-155B	PS8326-30	PS8326-100	PS8327-30	PS8327-100	PS8329-30	PS8329-100	PS8330-30	PS8330-100	PS8331-30	<b>S8331</b>	PS8333-30	PS8333-100

Table 9. Availability of Sediment Trap Data for 1983

0 Trap Name	1 s/N	2 Mooring	3 Pig- ments	4 Carbon	5 Size Frac.	6 TM	7 Org.	8 Page
PS833-50	26	PS833	x					56
PS833-50B		PS833	X					57
PS833-100	27	PS833	X					58
PS834-50	28	PS834	x					59
PS834-100	29	PS834	X		v			60 61
PS835-31	14	PS835	x		x	*		61 95
PS835-31	1 =	20025	v			x		62
PS835-50 PS835-50	15	PS835	X			x		95
PS835-50B	16	PS835	x					63
PS835-50C	17	PS835	x					64
PS835-100	18	PS835	x		x			65
PS835-100	10	1 5000				x		95
PS835-148	20	PS835	x					66
PS835-148						x		95
PS835-214	23	PS835	x					67
PS835-214						x		95
PS836-50	24	PS836	x					68
PS836-100	25	PS836	x					69
PS838-50	2	PS838	X					70
PS838-50						х		95
PS838-100	6	PS838	x					71
PS838-100						X		96
PS838-166	7	PS838	x					72
PS838-166						x		95 73
PS8310-50	11	PS8310	x					73
PS8310-50	10	D00310	•			x		95 74
PS8310-100 PS8310-100	12	PS8310	x			x		95
	13	DC9210	x			_		75
PS8310-170 PS8310-170	13	PS8310	A.			x		95
PS8312-88	30	PS8312	x					76
PS8312-88	30	FDOJIZ				x		95
PS8324-30	31	PS8324	x					77
PS8324-30B	32	PS8324	x					78
PS8324-90	33	PS8324	X					79
PS8324~90B	34	PS8324	x					80
PS8334-155	35	PS8324	x					81
PS8324-155B	36	PS8324	x					82
PS8326-30	17	PS8326	x					83
PS8326-30							x	97
PS8326-100	15	PS8326	X					84
PS8326~100							x	97
PS8327-30	14	PS8327	X		x			85
PS8327-30	4.0		••				x	97
PS8327-100	18	PS8327	x		X		***	86
PS8327-100	1.0	ma0200	**				X	97
PS8329-30 PS8329-30	16	PS8329	x				x	87 97
PS8329-30 PS8329-100	20	PS8329	v				<b>.</b>	88
PS8329-100 PS8329-100	20	F 50 3 2 3	X				x	97
PS8330-30	2	PS8330	X		x			89
PS8330-30	L	E 20220	A				x	97
PS8330-30	6	PS8330	x		x			90
PS8330-100	•						==	
PS8330-100 PS8331-30	7	PS8331	x				x	97
PS8331-30	•	E 2022T	A				v	91 97
PS8331-100	11	PS8331	x				X	97
PS8331-100		100001	A				x	92 97
PS8333-30	13	PS8333	x				Δ.	93
PS8333-100	23	PS8333	x					94
PS8333-100	_		_ <del>_</del>				x	97

## 3.4 1984 Sampling Year

The sampling at 30 and 100 m at the locations off Point No Point (PS8402), in central main basin (PS8403) and off Three Tree Point (PS8405) that began in 1983 continued in 1984 between 1 December 1983 and 30 March 1984 (Table 10) and between 5 April 1984 and 23 July 1984 (PS8413, PS8414 and PS8415, respectively). During the 1 December 1983 to 30 March 1984 period, sediment traps were also deployed at 30, 160 and 195 m at the central main basin location (PS8403), at 32 and 88 m in the Strait of Juan de Fuca (PS8401), at 108 and 165 m in Whidbey Basin (PS8407) and at 30 and 100 m off Browns Point (PS8406).

The trace organics data (Table 11) is interpreted in Murphy et al. (1988).

Table 10. Sediment Trap Location and Sampling Data for 1984

10 Comments			of Double		
	84	0 0 0 0 4 4 4 4	0 0 0 0 0 4 4 4 4	8 8 8 8 8 8 4 4 4 4 4	84 16:00 84 16:00 84 16:00 84 16:00 84 16:00 84 16:00
9 Stop: Date/	1				30 MAR 23 JUL 23 JUL 23 JUL 23 JUL 23 JUL 23 JUL 23 JUL
8 Duration Cylinder	288 288	2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	7 7 7 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8 8	888888888888888888888888888888888888888	0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0 0
7 Start: Date/	1 DEC 83	1 DEC 83 1 DEC 83 1 DEC 83 1 DEC 83	1 DEC 83 1 DEC 83 1 DEC 83 1 DEC 83	1 DEC 83 1 DEC 83 1 DEC 83 1 DEC 83 1 DEC 83	1 DEC 83 20:00 5 APR 84 16:00 5 APR 84 16:00 5 APR 84 16:00 5 APR 84 16:00 5 APR 84 16:00
6 Sample Depth	32	9 9 9 2 3 8 9 20 5 9	988 988 160 195	32 98 32 32 108	168 168 168 168 168 168 168 168 168 168
5 Water Depth	128	194 194 201 201	201 201 201 201	221 221 172 172 221	221 2200 2200 221 221 121
4 Long.	122 57.8 122 57.8			122 24.3 122 24.3 122 26.5 122 26.5 122 26.5	122 20.5 122 28.8 122 28.8 122 27.4 122 27.4 122 24.3
3 Lat.	8 13.	47 53.9 47 53.9 47 41.7 47 41.7	47 41.7 47 41.7 47 41.7	47 27.0 47 27.0 47 19.9 47 19.9	27.055.3 27.44.55.3 27.44.59.8 27.10.7 27.0
2 Mooring 3 Lat		PS8402 PS8402 PS8403 PS8403	PS8403 PS8403 PS8403 PS8403	PS8405 PS8405 PS8406 PS8406 PS8407	PS88407 PS88413 PS88413 PS88414 PS8414 PS8415
1 S/N	15 16	11. 13. 14. 14.	2 5 2 5 2 5 2 5 5 5 5 5 5 5 5 5 5 5 5 5	28 30 31 12	34 20 32 36 36
O Trap Name	PS8401-32 PS8401-88	PS8402-30 PS8402-98 PS8403-32 PS8403-50	PS8403-98 PS8403-98B PS8403-160 PS8403-195	PS8405-32 PS8405-98 PS8406-32 PS8406-98 PS8407-108	PS8407-165 PS8413-32 PS8413-98 PS8414-32 PS8414-98 PS8415-32 PS8415-32

Table 11. Availability of Sediment Trap Data for 1984

0 Trap Name	1 S/N	2 Mooring	3 Pig- ments	4 Carbon	5 Size Frac.	6 TM	7 Org.	8 Page
PS8401-32	15	PS8401	x					98
PS8401-32	1.7	10401	-			x		120
PS8401-32						_	×	122
PS8401-88	16	PS8401	x					99
PS8401-88						X		120
PS8401-88							x	122
PS8402-30	17	PS8402	x					100
PS8402-30						X		120
PS8402-98	18	PS8402	x					101
PS8402-98						X		120
PS8403-32	33	PS8403	X					102
PS8403-50	24	PS8403	X					103
PS8403-50						X		120
PS8403-98	25	PS8403	X					104
PS8403-98						X		120
PS8403-98B	25	PS8403	x					105
PS8403-160	26	PS8403	x					106
PS8403-160						X		120
PS8403-195	27	PS8403	x					107
PS8403-195						x		120
PS8405-32	28	PS8405	x					108
PS8405-32	20	D0040E	••			X		120
PS8405-98	29	PS8405	x			••		109
PS8405-98 PS8406-32	30	PS8406	•			X		120
PS8406-32	30	P38406	x			x		110 120
PS8406-32	31	PS8406	x					111
PS8406-98	31	F30400				x		120
PS8407-108	12	PS8407	x					112
PS8407-108		I DOTO	4			x		120
PS8407-108							x	122
PS8407-165	34	PS8407	x				-	113
PS8407-165			_			x		120
PS8407-165							x	122
PS8413-32	2	PS8413	x					114
PS8413-98	7	PS8413	x					115
PS8414-32	13	PS8414	X					116
PS8414-32							x	122
PS8414-98	20	PS8414	x					117
PS8414-98							x	122
PS8415-32	32	PS8415	X					118
PS8415-32							X	122
PS8415-98	36	PS8415	X					119

# 3.5 1985 Sampling Year

A single mooring (PS8509) was deployed in the central main Basin between 29 March 1985 and 22 June 1985 at depths of 30, 100 and 175 m (Table 12). A second sediment trap at 100 m (100s) had a cylinder duration time of 4 hours.

Sedimentation rates and Mn concentrations on settling particles (Table 13) are discussed in Cudaback et al. (1991).

			Table	12. Sedim	ent Trap	Location a	Table 12. Sediment Trap Location and Sampling Data for 1985	tor 1985		
O Trap Name	1 S/N	1 s/N 2 Mooring	3 Lat.	4 Long. 5 W	5 Water Depth	Water 6 Sample 7 Start: Depth Depth Date/	7 Start: Date/	8 Duration Cylinder	n 9 Stop: c Date/	10 Comments
PS8509-30 25 PS8509 47 42.0 1	25	PS8509	47 42.0	122 27.1 195	195	30	30 29 MAR 85 00:00	204	204 22 JUN 85 00:00	
PS8509-100	26	PS8509	47 42.0	122 27.1	195	100	29 Mar 85 00:00	204	22 JUN 85 00:00	
PS8509-177	27	PS8509	47 42.0	122 27.1	195	175	29 MAR 85 00:00	204	22 JUN 85 00:00	
PS9809-100S	11	PS8509	47 42.0	122 27.1	195	100	10 APR 85 09:00	4	12 APR 85 01:00	

Availability of Sediment Trap Data for 1985 Table 13.

O Trap Name	1 S/N	1 S/N 2 Mooring 3 Pig- 4 Carbon 5 Size 6 TM 7 Org. 8 Page ments Frac.	3 Pig- ments	ਹ ਵਾ	arbon	5 Size Frac.	MI 9	7 org.	8 Page
PS8509-30 25 PS8509 x 123	25	PS8509	 	i ! !	)           	· · · · · · · · · · · · · · · · · · ·	×		123
PS8509-100	56	PS8509							124
PS8509-175	27	PS8509							125
PS8509-175							×		127
PS8509-100S	11	PS8509							126
PS8509-100S							×		127

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