A Pilot Study for Ambient Toxicity Testing in Chesapeake Bay

Volume I Year 1 Report



A Pilot Study for Ambient Toxicity Testing in Chesapeake Bay

Volume I - Year I Report

April, 1991

Printed by the U.S. Environmental Protection Agency for the Chesapeake Bay Program

Prepared by:

Lenwood W. Hall, Jr. Michael C. Ziegenfuss Steven A. Fischer

University of Maryland Agricultural Experiment Station Wye Research and Education Center

> Raymond W. Alden, III Emily Deaver

Old Dominion University
College of Sciences
Applied Marine Research Laboratory

Jay Gooch Nikki DeBert-Hastings

University of Maryland Chesapeake Biological Laboratory

Produced in cooperation with the Maryland Department of Natural Resources,
Maryland Department of the Environment and
U.S. Environmental Protection Agency Chesapeake Bay Program Office

FOREWORD

This pilot study was designed to evaluate ambient toxicity in the Chesapeake Bay watershed by using a battery of water column, sediment and suborganismal tests. To complete this ambitious goal, of scientists from three Chesapeake Bay Research Laboratories were involved. Water column toxicity studies were directed by Lenwood W. Hall, Jr. of the University of Maryland's Agricultural Experiment Station, sediment toxicity tests were managed by Raymond W. Alden, III of Old Dominion University Applied Marine Research Laboratory, and suborganismal tests were directed by Jay Gooch of the University of Maryland's Chesapeake Biological Laboratory. This report summarizes data from the first year of The following government agencies were this two-year study. responsible for supporting and/or managing this research: Maryland (CB-90-001-005), Natural Resources Maryland Department of Department of Environment (182-C-MDE-91) and the U.S. Environmental Protection Agency (X-003554-01).

ABSTRACT

The link between toxic contaminants and biological effects in the ambient environment of the Chesapeake Bay watershed has become a major environmental issue in recent years. Evaluating potentially toxic conditions in natural habitats can only be accomplished by direct measurement of biological responses in the ambient environment. The primary goal of this ambient toxicity pilot study was to identify toxic areas in the Chesapeake Bay watershed by using a battery of concurrent water column, sediment and suborganismal toxicity tests. Ambient areas evaluated in the first year of this two-year pilot study were: suspected toxic locations in the Elizabeth River (one station) and Patapsco River (one station); a suspected contaminant-free area in the Wye River (one station) and critical habitat areas in the Potomac River (three freshwater and two saltwater stations). The Potomac River stations were located at Indian Head, Freestone Point, Possum Point, Morgantown and Dahlgren. A suite of inorganic and organic contaminants was evaluated in the water column and sediment from at least one grab sample collected from selected stations during water column and sediment tests. Standard water quality conditions were also evaluated in water and sediment from all stations.

The following water column tests were conducted in saltwater: 8-d sheepshead minnow, <u>Cyprinodon variegatus</u>, survival and growth tests; 8-d grass shrimp, <u>Palaemonetes pugio</u>, survival and growth tests and 8-d Eurytemora affinis life cycle test. In freshwater habitats of the Potomac River, 8-d fathead minnow, Pimephales promelas, larval survival and growth tests and 8-d Ceriodaphnia dubia survival and reproduction tests were conducted. Concurrent tests with salinity adjusted freshwater were conducted at the freshwater stations with E. affinis and P. pugio to compare these results with those from the two freshwater tests. The following sediment tests were conducted: 20-d juvenile and adult polychaete worm, Streblospio benedicti, survival and growth test; 20-d juvenile grass shrimp , P. pugio, survival and growth test and 20-d amphipod, Lepidactylus dytiscus, survival, growth and reburial Concurrent tests with salinity adjusted freshwater were test. conducted with the above three species to compare these results with those from the freshwater tests. Suborganismal tests were conducted to evaluate hepatic monooxygenase (EROD) activity in Fundulus heteroclitus exposed to test water and sediment from the saltwater stations. Livers were collected from Fundulus exposed to water and sediment and analyzed for induction of cytochrome P450 monooxygenase activity indicative of aromatic hydrocarbon exposure. In addition, feral Fundulus were collected where possible from near the test stations and analyzed for induced monooxygenase activity. White perch, Morone americana, and/or spot, Leiostomus xanthurus, were collected by trawling or hook and line in areas where Fundulus were not found. Livers from these fish species were also analyzed for induced monooxygenase activity.

The Elizabeth River was the most toxic area tested based on

the various biological indicators. The water column and sediment tests demonstrated the presence of toxic conditions at this station and the suborganismal tests suggested toxic conditions. Patapsco River station was also reported toxic as water column and sediment tests indicated adverse conditions to be present. Feral spot collected at this station had EROD hepatic activities that were similar to spot collected from the highly PAH-contaminated Elizabeth River station. The Wye River station was selected as a putative toxic-free control area. Water column tests did not suggest the presence of toxic conditions in this area but sediment tests demonstrated significant effects. Feral Fundulus from the Wye River had hepatic EROD activities which suggested possible planar aromatic hydrocarbon exposures. Both water column and sediment tests demonstrated toxic conditions at the Indian Head, Sediment tests showed Morgantown and Dahlgren stations. significant effects at the Freestone Point and Possum Point stations while no effects were reported at these stations from water column or suborganismal tests. The results discussed above clearly demonstrate the need for integrated water column, sediment and suborganismal testing because one type of test was not sufficient to maximize our ability to identify toxic conditions in the ambient environment of the Chesapeake Bay watershed.

A general ranking of sensitivity among tests within each test type (water column, sediment and suborganismal) showed the following results after one year of testing. Results from the water column tests demonstrated no significant ranking of

sensitivity among the three saltwater tests but rather supported the need for multispecies tests because different species displayed varying sensitivity to different types of contaminants. Both grass shrimp and sheepshead minnow tests demonstrated effects at two of the eight stations while E. affinis showed effects at only one station. A comparison of the two freshwater water column tests at three stations showed that the Ceriodaphnia test was more sensitive than the fathead minnow test. Results from the sediment tests clearly showed that the amphipod test was most sensitive as effects were reported at all stations. The polychaete worm test was the second most sensitive test and the grass shrimp test was least of among the various sensitivity ranking sensitive. A suborganismal tests is not appropriate until further research has been completed.

organic and inorganic contaminants data from the water column and sediment of the various stations provided supportive but not conclusive evidence to explain biological effects. The greatest number of contaminants at possibly toxic concentrations (three for water and fourteen for sediment) were reported at the Elizabeth River station, where all three types of tests demonstrated toxic conditions. Three to four metals at possibly toxic concentrations were found in the sediment at the Patapsco River, Indian Head, Freestone Point and Possum Point stations. Sediment toxicity was demonstrated at all of the stations with the most sensitive sediment toxicity test (amphipod test). This sediment test also showed toxicity at the Morgantown and Dahlgren stations where

metals at possibly toxic concentrations were not reported in the sediment. One or two metals at possibly toxic concentrations were reported in the water column at the following stations: Patapsco River, Wye River, Indian Head, Freestone Point and Morgantown. Significant effects from water column tests were reported at the Patapsco River, Indian Head, Morgantown and Dahlgren stations; effects were not reported at the Wye River, Possum Point or Freestone Point stations. Suborganismal tests suggested planar aromatic hydrocarbon exposure at the Patapsco River and Wye River stations but effects were not implicated at any of the five Potomac River stations.

ACKNOWLEDGEMENTS

We would like to acknowledge the following individuals or organizations for assisting in this study: John Sullivan and Deborah Palmer for technical assistance; Mary Hancock for typing; Irvin Maddox for the use of his boat; VERSAR, Inc. for contaminant analysis and Mel Knott for providing toxicity data on the Patapsco River. Michael Hirshfield, Richard Batiuk and Mary Jo Garreis are acknowledged for their comments on the study design.

TABLE OF CONTENTS

		<u>Page</u>
	Foreword	i
	Abstract	ii
	Acknowledgements	vii
	Table of Contents	viii
1.	Introduction	1
2.	Objectives	4
3.	Methods	8
	3.1 Study Areas 3.1.1 Elizabeth River Station	8 8
	3.1.2 Patapsco River Station	11 12
	3.1.3 Wye River Station 3.1.4 Potomac River - Indian Head Station	12
	3.1.5 Potomac River - Freestone Point Station	13
	3.1.6 Potomac River - Possum Point Station	14
	3.1.7 Potomac River - Morgantown Station	14
	3.1.8 Potomac River - Dahlgren Station	15
	3.2 Water Column Tests	15
	3.2.1 General Description	15 16
	3.2.2 Culture and Maintenance for Test Species 3.2.2.1 Grass shrimp	18
	3.2.2.1 Grass Shrimp 3.2.2.2 Copepod - <u>E</u> . <u>affinis</u>	18
	3.2.2.3 Sheepshead minnow	19
	3.2.2.4 Fathead minnow	20
	3.2.2.5 <u>Ceriodaphnia</u>	20
	3.2.3 Reference Toxicant Tests	21 22
	3.2.3.1 Grass shrimp	23
	3.2.3.2 Copepod - <u>E</u> . <u>affinis</u> 3.2.3.3 Sheepshead minnow	23
	3.2.3.4 Fathead minnow	24
	3.2.3.5 Ceriodaphnia	24
	3.2.4 Sample Collection, Handling and Storage	25
	3.2.5 Test Procedures	26 26
	3.2.5.1 Grass Shrimp	27
	3.2.5.2 Copepod - <u>E</u> . <u>affinis</u> 3.2.5.3 Sheepshead minnow	29
	3.2.5.4 Fathead minnow	31
	3.2.5.5 Ceriodaphnia	32
	3.2.5.6 Fundulus heteroclitus	34
	3.2.6 Contaminant and Water Quality Evaluations	34

			<u>Page</u>
	3.3	Sediment Tests 3.3.1 General Description	38 38
	-	3.3.2 Collection, Handling, Maintenance and	42
		Culture of Test Species 3.3.2.1 Grass shrimp - Palaemonetes pugio	42
		3.3.2.2 Amphipod - <u>Lepidactylus dytiscus</u> 3.3.2.3 Polychaete worm - <u>Streblospio</u>	42
		<u>benedicti</u>	43
		3.3.3 Reference Toxicant Tests	43
		3.3.4 Sample Collection, Handling and Storage	44
		3.3.5 Test Procedures	46
		3.3.5.1 Grass shrimp - <u>Palaemonetes pugio</u> 3.3.5.2 Amphipod - <u>Lepidactylus dytiscus</u> 3.3.5.3 Polychaete worm - <u>Streblospio</u>	48 48
		benedicti 3.3.6 Contaminant and Sediment Quality	49
		Evaluations 3.3.6.1 Semi-volatile Organic Analysis:	49
		Sediment	49
		3.3.6.2 Pesticide Analysis: Sediment	53
		3.3.6.3 Acid Volatile Sulfides	53
		3.3.6.4 Total Organic Carbon	56
		3.3.6.5 Ammonia and Nitrite 3.3.6.6 Metals	56 57
	3.4	Suborganismal Tests	5 7
		3.4.1 Fundulus - Water and Sediment Exposures	57
		3.4.2 Collection of Feral Fishes	58
		3.4.3 Sampling of Organisms from Water and	
		Sediment Toxicity Test	59
		3.4.3.1 <u>Fundulus</u>	59
		3.4.3.2 Larval Fish and Invertebrates	60
		3.4.3.3 Biochemical Methods	60
		3.4.3.4 Stress Proteins	61
4.	Resu	nlts	63
	4.1		63
		4.1.1 Toxicity Data	63
		4.1.2 Contaminants Data	71
		4.1.3 Water Quality Data 4.1.4 Reference Toxicant Data	74 74
	4.2	Sediment Tests	77
		4.2.1 Toxicity Data	77
		4.2.2 Contaminants Data	85
		4.2.3 Pore Water Data	89
		4.2.4 Reference Toxicant Data	91

			Page	
	4.3 Suborganismal Tests 4.3.1 Cytochrome P-450 Induction 4.3.1.1 Fundulus - Water and Sediment			
5.	Disc	Exposures 4.3.1.2 Larval Sheepshead and Fathead Minnows from Water Exposures 4.3.1.3 Feral Fish 4.3.1.4 Ongoing Activities ussion	93 96 97 99 101	
J .	5.1	Elizabeth River Station	101	
	5.2	Patapsco River Station	107	
	5.3	Wye River Station	111	
	5.4	Potomac River - Indian Head Station	114	
	5.5	Potomac River - Freestone Point Station	115	
	5.6	Potomac River - Possum Point Station	117	
	5.7	Potomac River - Morgantown Station	118	
	5.8	Potomac River - Dahlgren Station	120	
6.	Cond	clusions	122	
7.	Ref	erences	127	
	App	endices		
	A.	Suborganismal sampling for various fish species.		
	в.	Water quality data collected at 1-h intervals with datasonde units during the Potomac River experiments.		
	c.	Water quality conditions reported in test containers during all water column tests.		
	D.	Organic analysis data for Possum Point, Freestone Point, Dahlgren, Elizabeth River and Nansemond River sediment.		
	Ε.	Pesticide analysis data for Possum Point, Freestone Point, Dahlgren, Elizabeth River and Nansemond River sediment.		

SECTION 1

INTRODUCTION

In recent years there has been concern about the link between adverse biological effects and the presence of toxic contaminants and adverse water quality conditions in critical Chesapeake Bay habitat areas. Information on the loading of toxic contaminants is not sufficient to determine the biological effects of complex mixtures of contaminants derived from sources such as atmospheric deposition, groundwater contamination, multiple point source effluents, non-point source runoff from agricultural and urban land and release of toxic contaminants from sediments. Evaluating potentially toxic conditions in natural habitats can only be accomplished by direct measurement of biological responses in the ambient aquatic environment. Numerous investigators have reported toxic conditions in ambient waters of the Chesapeake Bay watershed (Alden et. al., 1988; Bender and Huggett, 1987; Hall et al., 1985; Hall et al., 1989; Klauda et al., 1988; Wright et al., 1988).

Ambient toxicity tests are designed to evaluate toxicity in the ambient aquatic environments (areas outside of any mixing zones) by using biological responses (survival, growth, reproduction, etc.) of test species. Ambient toxicity tests are conducted less frequently when compared with the more traditional effluent toxicity tests. Effluent toxicity tests are used to establish effluent discharge rates that minimize effects on

organisms in the receiving system (U.S. EPA., 1983; U.S. EPA., 1985a). Ambient toxicity tests are designed to detect toxic conditions in the ambient environment (areas outside of mixing zones) by using responses of aquatic biota. Toxic conditions in the ambient environment may result from point or non-point sources or by combinations of effluents that are not toxic individually.

Toxicity can be defined in a number of ways. toxicity testing protocols used for regulatory purposes focus primarily on toxicity as defined by effects on survival, reproduction and growth of test species. In recent years, increases in our knowledge of how chemicals affect biochemical systems (i.e., mechanisms of toxicity) has lead to the development of tests, or more appropriately, to the use of biochemical effects measurements (suborganismal tests) to detect sublethal levels of exposure in laboratory and field experiments. In this study, we have addressed ambient toxicity testing in a comprehensive manner by using concurrent water column, sediment and suborganismal tests in conjunction with water quality and contaminant analysis in both water and sediment. Both ambient water and sediment were collected and transported back to the laboratory for testing. This battery of tests will maximize our ability to detect "ambient toxicity" in Chesapeake Bay waters during this pilot study.

The Chesapeake Bay Basinwide Toxics Reduction Strategy has a commitment to develop and implement a plan for Baywide assessment and monitoring of the effects of toxic substances, within natural habitats, on selected commercially, recreationally and ecologically

important species of living resources (CEC, 1988a). This commitment is consistent with recommendations of the Chesapeake Bay Living Resource Monitoring Plan (CEC, 1988b). In July of 1989 an Ambient Toxicity Assessment Workshop was held in Annapolis, Maryland, to provide a forum on how to use biological indicators to monitor the effects of toxic contaminants in Chesapeake Bay habitats to living resources. This pilot study was designed by using various recommendations of the workshop.

SECTION 2

OBJECTIVES

The objectives of year one of this two-year pilot study were to: (1) develop a survey program to broadly assess ambient toxicity of living resource habitats for the purpose of identifying defined regions where ambient toxicity levels warrant further investigation of effects on living resources; (2) assess the feasibility and utility of such a survey through a pilot study; (3) field test existing standardized, directly modified or recently developed water column, sediment and suborganismal toxicity test methods for use in ambient toxicity testing and determine the relative sensitivity of these three test methods and (4) identify long term test methods development or follow up survey design testing needs (if any) to support Baywide assessment of ambient toxicity. Toxicity in natural habitat areas was evaluated by using water column toxicity tests (Lenwood Hall), sediment toxicity tests (Ray Alden) and suborganismal evaluations (Jay Gooch).

Test areas included polluted areas (Elizabeth River and Patapsco River), critical habitat areas with potentially toxic conditions (Potomac River), and a presumed contaminant-free area (Wye River). One station was located in the Elizabeth River, Patapsco River and Wye River, and five stations were located in the Potomac River (3 freshwater and 2 saltwater). All eight stations were evaluated once in 1990 (August 13 - October 2). A suite of

inorganic and organic contaminants was evaluated in the water column and sediment for selected station during testing.

Water column, sediment and suborganismal tests were conducted concurrently. The following saltwater water column tests were conducted: 8-d sheepshead minnow, Cyrinodon variegatus, survival and growth test; 8-d grass shrimp, Palaemonetes pugio, survival and growth test and 8-d copepod, Eurytemora affinis, life cycle test. In freshwater habitats, 8-d fathead minnow, Pimephales promelas, larval survival and growth test and 8-d Ceriodaphnia dubia survival and reproduction tests were conducted. Concurrent tests with salinity adjusted freshwater were conducted at the freshwater stations with E. affinis and P. pugio to compare these results with those from the two freshwater tests.

The following saltwater sediment toxicity tests were conducted: 20-d juvenile <u>Streblospio benedicti</u> (polychaete worm) survival and growth test; 20-d juvenile grass shrimp (P. pugio) survival and growth test and 20-d amphipod (<u>Lepidactylus dytiscus</u>) survival, growth and reburial test.

Several types of samples were collected for suborganismal testing and many of the analyses are still ongoing. Our efforts to date have been focused on evaluating hepatic monooxygenase activity in feral and laboratory-exposed fishes. Livers were collected from Fundulus heteroclitus exposed to test sediment and water and analyzed for induction of cytochrome P450 monooxygenase activity indicative of aromatic hydrocarbon exposure. In addition, feral Fundulus were collected, where possible, from near the test sites

and analyzed for induced monooxygenase activity. For stations where we were unable to collect resident <u>Fundulus</u> (inappropriate habitat or salinity), we collected white perch (<u>Morone americana</u>) and/or spot (<u>Leiostomus xanthurus</u>) by trawling or hook and line. Livers from these fish species were also analyzed for induced monooxygenase activity.

Samples of animals were also collected from various water and sediment toxicity tests and preserved in liquid nitrogen. These samples will be analyzed in a logical progression. We collected samples from the grass shrimp (P. pugio) and amphipod (L. dytiscus) sediment tests for analysis of stress protein responses. We also collected samples from the invertebrate water column tests (grass shrimp and Eurytemora) and samples of surviving sheepshead and fathead minnow larvae for stress protein evaluations. We will also be evaluating monooxygenase induction in the larval fish samples.

An important goal of this integrated water column, sediment and suborganismal testing was to use this battery of tests as a screening tool to identify toxic areas in ambient waters of the Chesapeake Bay watershed. Results from this study will be used to determine if toxicity exists in ambient areas. Inorganic contaminants, organic contaminants and water quality conditions (temperature, dissolved oxygen, pH, salinity and conductivity) were evaluated during these tests. However, the limited analysis of these contaminants and water quality conditions was not intended to provide conclusive evidence on what factors caused toxicity (if reported). In addition, this study was not designed to identify

sources of contaminants if they were reported. The use of these data for regulatory processes would only be warranted after more comprehensive studies were designed to evaluate causes and sources of toxicity. Data collected during this study would, however, be useful for identifying potentially toxic areas that would be of interest to regulatory agencies.

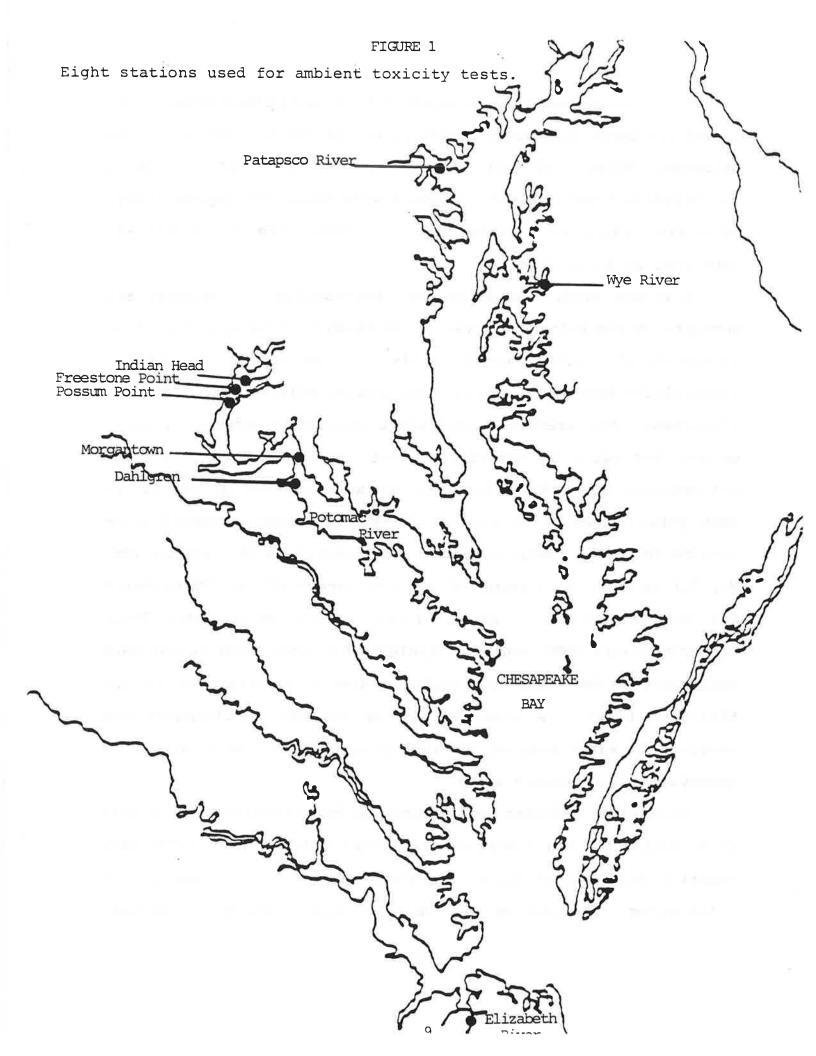
SECTION 3

METHODS

3.1 Study Areas

Three types of areas were selected for pilot ambient toxicity tests: polluted areas in the Elizabeth River and Patapsco River; a relatively toxic-free area in the Wye River; critical habitat areas in the Potomac River with potentially toxic conditions (Figure 1). The polluted sites in the Elizabeth River and Patapsco River were selected to allow field testing of existing test methods for ambient toxicity testing. Reduced survival and other sublethal effects would be likely with some of the test species. River was selected because it is a relatively clean area that should not have toxic conditions. Conducting ambient toxicity tests in the Wye River will provide data to demonstrate that the proposed test methods will demonstrate adequate survival or no sublethal effects by test species in toxic-free water. The Potomac River was selected for testing because it offers gradients of salinity, contamination and habitat similar to those of the whole Bay. Another reason for selecting the Potomac River is that the geography and numerous resources of the river are shared by three states. A brief description of each study site and rationale for selection are presented below.

3.1.1 Elizabeth River Station



This station was located adjacent to Atlantic Wood Industries, a wood treatment (creosote) industry, on the Southern Branch of the Elizabeth River (36° 48' 29" N x 76° 17' 36" W) in a mesohaline/polyhaline area. Samples were collected approximately 30 m from shore at a depth of 6 m. This site is on the EPA Superfund National Priority List.

A recent study evaluating the distribution of toxicants and mutagens in the Elizabeth River in relation to land use activities (Alden et al., 1988) reported this site as having the highest quantitative ranking for overall biological effects among 27 sites For example, significant cumulative effects (acute, chronic and sublethal effects) of contaminant extracts from water collected at this site on fish embryos were observed in >50% of the The site also had the highest quantitative test populations. ranking for water chemistry data. Concentrations of 3 metals (Cu, Ni, Zn) in water were reported to be in excess of the EPA criteria Total phosphorus, Total for the protection of marine life. Suspended Solid (TSS) and Total Kjeldahl Nitrogen (TKN) levels were noticeably higher than the ambient water concentrations in the Elizabeth River. A wide variety of organic contaminants was observed in water samples, including numerous alkanes, alkylated aromatics and propanoic acids.

Studies of sediment contaminants in the vicinity of this site (i.e. within 0.5 km) (Ewing et al., 1989; Alden et al., 1990) have reported potentially toxic concentrations of Cd (>3 mg/kg), Cr (>100 mg/kg), Cu (>300 mg/kg), Pb (>400 mg/kg) and Hg (>3 mg/kg).

Concentrations of organic contaminants in sediments were generally high in this portion of the Elizabeth River (e.g. total PNAHs approximately 150 mg/kg).

A study conducted by the Virginia Institute of Marine Sciences (VIMS) reported high concentrations (i.e. highest in the Southern Branch) of total aromatic organic compounds in the tissues of Callinectus (3,200 ppb in muscle, 13,000 ppb in hepatopancreas) collected from this site (Greeves, 1990). Other VIMS researchers have observed an exceptionally high incidence of neoplastic liver lesions in Fundulus from this site (Vogelbein et al., 1990).

Bioassessments conducted by Old Dominion University Applied Marine Research Laboratory in early summer, 1990, showed significant toxic effects in tests of sediment pore water (Microtox and sea urchin fertilization tests) and whole sediments (amphipod tests) from this site (Emily Deaver, personal observation). This station, located in a documented toxic area of the Elizabeth River, was selected to allow field testing of our tests in an area where effects were likely.

3.1.2 Patapsco River Station

This station was located adjacent to the Bethlehem Steel Corporation at Sparrows Point on the Patapsco River near Bear Creek (39° 14' N x 76° 29.7' W). Samples were collected approximately 75 m from shore at a depth of 2 m. This site was located in a mesohaline area outside the mixing zone of Bethlehem Steel's largest discharge. Effluent toxicity tests conducted by the

University of Maryland's Agricultural Experiment Station have reported chronic effects on Bethlehem Steel's effluent from both Ceriodaphnia and fathead minnow tests (Mel Knott, personal communication). Various potentially toxic contaminants such as zinc, chromium, lead-free cyanide and tetrachloroethane have been reported in effluents from the Bethlehem Steel facility.

This station was selected to allow field testing of our methods in an area where toxic conditions may be found. Based on our limited background data, the ambient environment at this station should be less toxic than the Elizabeth River station.

3.1.3 Wye River Station

This station was located in a mesohaline area on the Wye River at Wye Narrows above the Manor House (38° 53.2' N x 76° 1.9' W). Samples were collected approximately 75 m from shore at a depth of 2 m. There are no known point sources of contaminants in this area and non-point runoff is minimal due to the vegetation cover and hardwoods in the area. This area supports healthy populations of various Chesapeake Bay species. A preliminary sample of water from this area did not indicate any potentially toxic contaminants. This station was selected as a control area where toxic conditions were not suspected.

3.1.4 Potomac River - Indian Head Station

This station was located in a freshwater area on the Maryland side of the Potomac River near the Naval Ordnance Station (NOS) at

Indian Head, Maryland (38° 36' N x 77° 11' W). Samples were collected approximately 125 m from shore at a depth of 4 m. The NOS facility has 51 industrial outfalls and one municipal outfall discharging into the Potomac River. In recent screening toxicity tests with freshwater species, several of the outfalls were toxic. We have no documented evidence to demonstrate that the NOS facility is causing adverse effects to aquatic life in the ambient areas of the Potomac River. However, the station near this facility is a representative ambient area of the Potomac River and provides a good location for the security of our equipment during testing (datasonde units).

3.1.5 Potomac River - Freestone Point

This station was located in a freshwater area on the Virginia side of the Potomac River near the mouth of Nebesco Creek (38° 35.2' N x 77° 14.7' W). Samples were collected approximately 100 m from shore at a depth of 4 m. This station was near a station previously used for in-situ striped bass studies in 1986 (Hall et al., 1987a). The following metals were reported at this station in 1986: Cd (15 ug/L); Cr (110 ug/L) and Cu (47 ug/L). All of these concentrations exceed the U.S. EPA acute freshwater water quality criteria for these metals (U.S. EPA, 1987a). Higher than ambient pH conditions (> 9) were also reported at this station during the previously mentioned study. Survival of both striped bass prolarvae and yearlings was also low at this station during the insitu studies in 1986. This station was selected

representative area of the Potomac River where biological effects were possible.

3.1.6 Potomac River - Possum Point

This station was located in a freshwater area on the Virginia side of the Potomac River near Possum Point (38° 32' N x 77° 16.5' Samples were collected approximately 300 m from shore at a depth of 4 m. We conducted contaminant monitoring at this station during previous in-situ striped bass studies and reported the following concentrations of metals from one grab sample in 1990: Cd (4.7 ug/L), Cr (150 ug/L), Cu (37 ug/L), Ni (281 ug/L), Pb (14 ug/L) and Zn (270 ug/L) (Hall, personal communication). All of these metal concentrations exceed the U.S. EPA water quality criteria for freshwater (U.S. EPA, 1987a). Mortality of striped bass prolarvae was also high at this station during in-situ studies In 1989, above-ambient concentrations of arsenic (12 ug/L) and chromium (29 ug/L) were reported at this station (Hall et al., 1990). Significant mortality of striped bass prolarvae and yearlings was also reported during in-situ tests at this station. Previous potentially toxic contaminant conditions reported at this station during the spring of 1989 and 1990 were reasons for selecting this station.

3.1.7 Potomac River - Morgantown Station

This station was located in a typical ambient mesohaline area on the Maryland side of the Potomac River near the Harry Nice

Memorial Bridge (38° 21.8' N x 77° 59.5' W). Samples were collected approximately 200 m from shore at a depth of 3 m. TBT concentrations of 20 and 24 ug/L were detected at this station during a tributyltin (TBT) monitoring study in 1985 and 1986 (Hall et al., 1987b). These TBT concentrations exceed the U.S. EPA provisional water quality criteria of 10 ng/L (U.S. EPA, 1987b). There are no other background contaminants data for this station.

3.1.8 Potomac River - Dahlgren Station

This station was located in a typical ambient mesohaline area on the Virginia side of the Potomac River at the mouth of Machodoc Creek near the Naval Surface Warfare Center at Dahlgren, Virginia (38° 19' N x 77° 2' W). Samples were collected approximately 300 m from shore at a depth of 2 m. The Dahlgren facility has six discharge points in this area. This is a typical ambient mesohaline area of the Potomac River. Water quality and contaminant problems have not been reported in this area.

3.2 Water Column Toxicity Tests

3.2.1 General Description

The following estuarine tests were conducted with water collected at the Elizabeth River, Patapsco River, Wye River and Potomac River (Morgantown and Dahlgren stations): 8-d sheepshead minnow, Cyprindodon variegatus, survival and growth test; 8-d grass shrimp, Palaemonetes pugio, survival and growth test and 8-d Eurytemora affinis life cycle test. Three stations in the Potomac

River (Indian Head, Freestone Point and Possum Point) were located in freshwater areas. For these stations, 8-d fathead minnow, Pimephales promelas, larval survival and growth tests and 8-d Ceriodaphnia dubia survival and reproduction tests were performed. In addition to the two freshwater tests, estuarine tests with E. affinis and grass shrimp were conducted with salinity adjusted freshwater at these three sites. These estuarine tests were conducted concurrently with the two freshwater tests so results could be compared (i.e., significant biological effects or no significant biological effects). Control water used for freshwater tests was deep-well groundwater from our laboratory. Artificial seawater made with HW-MARINEMIX and deep-well groundwater was used as the control water for saltwater tests (15 °/00).

Each of the eight stations were tested one time during 1990 (August - October) with the various test species described above. Reference toxicant tests described later in this methods section were conducted at least once with each test species in 1990. The total number, type and location of tests conducted in 1990 are presented in Table 1. The Elizabeth River, Patapsco River and Wye River tests were conducted on 8/14/90 - 8/21/90. The Potomac River tests were performed on 9/24/90 - 10/2/90. A second test of Ceriodaphnia tests were conducted at the three freshwater Potomac River stations on 10/26/90 - 11/3/90.

3.2.2 <u>Culture or Maintenance Procedures for Test Species</u> The following subsections provide a brief description of culture or

The following abbreviations are for test species: C.v. (Cyprinodon variegatus) P.p. (Palaemonetes pugio), (Eurytemora affinis), P.pr. (Pimephales promelas), C.d. (Ceriodaphnia dubia). Water column toxicity tests conducted in 1990. used for test species: E.a. Table 1.

E.a.1			XXX		r	×	fac
P.p¹			XXX	T(*):	ć	××	
C.d.			XXX		(X ₂	×
Species P.pr.			XXX			× ₂ ×	×
ਜ਼ ਰ	×	×		XX	×	X ₂	×
P.p.	×	×		XX	×	X ₂	×
C. v.	×	×		XX	×	X ₂	×
Stations or Test Conditions (Number of Each)	Patapsco River (1)	Elizabeth River (1)	Potomac River Freshwater(3)	Potomac River Estuarine(2)	Wye River (1)	Control Water Column (1)	Water Column Ref. Tox. (1)

1 Salinity adjusted freshwater
2 Controls were used for each test

maintenance procedures for the five test species. Culture procedures for all test species except grass shrimp and \underline{E} . affinis are described in detail in Fisher et al. (1988).

3.2.2.1 Grass Shrimp

All grass shrimp used for reference toxicant or ambient toxicity tests were obtained from laboratory brood stock cultures at SP Engineering and Technology (Salem, MA). Larvae (< 7-d old) were held for 1 week in a 150 L fiberglass trough or 75 L aquaria at 15 ppt salinity and 25 C. A single airstone (medium trickle) was placed in the middle of the trough or aquaria to provide aeration and circulation. Temperature, salinity, dissolved oxygen, pH and ammonia were monitored daily prior to water renewal. Artificial seawater made with HW-MARINEMIX was used to exchange one-third of the water volume daily (HW-MARINEMIX, Wiegandt GMBH & Co., W. Germany). The bottom of the holding tank was siphoned daily to remove decaying organic matter. Larval grass shrimp were fed Artemia nauplii ad libitum one time per day following the water exchange. A 12-h L:12-h D cycle was maintained during the holding period.

3.2.2.2 Copepod (E. affinis)

Initial cultures of \underline{E} . affinis were obtained from established cultures at University of Maryland Chesapeake Biological Laboratory at Solomons Island, MD. Cultures were then maintained in our laboratory. Cultures were held in 10 L glass aquaria containing

synthetic seawater (HW-MARINEMIX) adjusted to 15 ppt salinity and 25 C. Stocking density was approximately 10³ organisms /L. Photoperiod was maintained at 12-h L:12-h D using indirect lighting. E. affinis were fed a phytoplankton diet consisting of Thalassiosira fluviatilis and Isochrysis galbana. Copepods were fed equal volumes of phytoplankton, in log-phase of growth, three times a week, to maintain a density of 10⁴ T. fluviatilis cells/ml and 10⁵ I. galbana cells/ml within the culture aquaria. The algal stocks were grown in 0.2um filtered well water adjusted to 15 ppt with synthetic sea salts using a standard culture procedure for marine algae (Guillard, 1975).

3.2.2.3 Sheepshead Minnows

Adult sheepshead minnows cultured in our laboratory were fed Tetramin® flake food ad libitum daily. Eggs obtained from these established cultures were used for the first set of experiments. Eggs were aerated vigorously in one liter glass beakers (150 eggs/L) containing synthetic seawater (HW-MARINEMIX) adjusted to 10 ppt salinity and 25C. Culture temperature and photoperiod (12-h L:12-h D) were controlled in biological incubators (Percival Manufacturing Co.). Dead eggs were removed from the beakers daily prior to 50 % culture water changes. Salinity was gradually adjusted to 15 ppt over two days to facilitate a uniform hatching rate. Twenty-four hours prior to starting a test, all previously hatched sheepshead larvae were separated from unhatched eggs and placed in a separate beaker. Only those larvae which hatched

within the twenty-four hours prior to starting the test were used to conduct the reference toxicant or ambient toxicity tests. Due to low hatching rate in our cultures, eggs were obtained from the Aquatic Bioassay Lab in Baton Rouge, LA, for the second set of experiments. The same procedures described above were used.

3.2.2.4 Fathead Minnows

Adult fathead minnows cultured in our laboratory were fed Tetramin® flake food ad libitum daily. Eggs obtained from these established cultures were used for our experiments. Eggs attached to a PVC substrate were placed in a five liter polyethylene container filled with 0.2 um filtered deep well water. The eggs were held at 25 C with moderate aeration. Twenty-four hours prior to tests, all previously hatched larvae were separated from unhatched eggs and collected in a separate beaker. Only those larvae which hatched within the 24 hours prior to starting the test were used to conduct the test.

3.2.2.5 Ceriodaphnia

Adult <u>Ceriodaphnia</u> were obtained from stock cultures at our laboratory. Organisms were cultured in one liter glass beakers containing 900 mL of 0.2 um filtered well water. The cultures were covered with clear plastic panels and gently aerated. The organisms were maintained at 25 C and 16-h L:8-h D photoperiod. Ceriodaphnia were fed a mixture of CEROPHYL (Sigma Chemical Co.) and <u>Selenastrum capricornutum</u>. The CEROPHYL was prepared as

previously described (Fisher et al., 1988) and fed at a rate providing approximately 60 ug/mL final concentration. Selenastrum capricornutum was cultured according to the methods described by the U.S. Environmental Protection Agency (U.S. EPA, 1985b) and fed at a rate to provide approximately 7 x 10⁵ cells/mL final concentration. Cultures were fed daily to maintain the organisms in optimum condition so as to provide maximum reproduction in the toxicity tests.

Four hours prior to starting the reference toxicant test, adult Ceriodaphnia were isolated in 250 ml glass beakers containing 60 mL of culture water (approximately 20 adults per beaker). After 4 h, neonates were collected by removing adults with a 500 um mesh Nitex screen. Reference toxicant and ambient toxicity tests were conducted on neonates that had hatched within 4 h prior to starting the tests.

3.2.3 Reference Toxicant Tests

A 48-h static cadmium chloride reference toxicant test was conducted with each test species in 1990. A reference toxicant is used to establish the relative health and sensitivity of the test organisms. Cadmium chloride was selected as the reference toxicant because we have a two-year data base with this chemical for all of the proposed test species except <u>E</u>. <u>affinis</u>. A complete description of rationale and use of reference toxicant tests is described in Fisher et al., 1988.

Cadmium concentrations were made from a common stock solution

of cadmium chloride (Sigma Chemical Co.). A 1 mg/mL (1.6308 mg/mL CdCl₂) cadmium stock solution was created by dissolving 407.7 mg CdCl₂ in 250 mLs deionized water. The stock solution was diluted to final concentrations by pipetting stock solutions into 2 L glass beakers containing filtered artificial seawater (15 ppt) diluent for saltwater tests. Deep well water was used for freshwater tests. Approximately 20 mL of each final test concentration was acidified with 100 ul ultrex nitric acid for future Atomic Absorption analysis.

Survival of test organisms was monitored at each test condition at both 24 and 48 hours (except <u>E</u>. <u>affinis</u> at 24 h). An LC50 value and 95 % confidence interval were calculated at 24 and 48 h using the moving-average and probit methods (Stephan, 1978). Temperature, salinity, dissolved oxygen, and pH were measured in each beaker at both 24 and 48 hours (except <u>E</u>. <u>affinis</u>). Specific details on methods used for each species are presented below.

3.2.3.1 Grass Shrimp

A 48-h cadmium chloride reference toxicant test was conducted with 18-d old larvae at the following nominal test concentrations (0, 0.18, 0.32, 0.56, 1.0, 1.8, 3.2 and 5.6 mg/L Cd). One replicate consisted of four larvae per 600 ml glass beaker containing 300 ml of test solution. Four replicates were conducted per test condition. Larvae were placed in individual Nitex chambers (0.202 mm mesh) in each beaker. We also exposed groups of 5 larvae per beaker in two additional beakers at 0.56 mg/L cadmium

to examine possible cannibalistic behavior.

3.2.3.2 Copepod (E. affinis)

A 48-h cadmium chloride reference toxicity test was conducted at the following nominal cadmium concentrations: 0, 0.032, 0.100, 0.180, 0.320, 0.560, and 1.000 mg/L. Ten to fifteen nauplii (< 24h old) were placed in replicate 150 ml glass beakers for each test condition. One hundred mls of test solution was used for each condition. A testing chamber was suspended within each beaker to The chambers were constructed from 3.8 cm hold the organisms. diameter polycarbonate rigid tubing cut to a length of 5.0 cm which provided a 40 ml volume when suspended in the beaker. The bottom of the chamber was covered with 53 um mesh Nitex screen. were counted by drawing small aliquots of nauplii (<24-h old) and culture water into a wide-bore, fire-polished glass pipette and examined under a dissecting microscope (15x magnification). number of nauplii introduced and the corresponding test chamber were recorded. Test beakers were placed in a biological incubator to maintain constant temperature (25 C) and photoperiod (12-h L:12h D). Mortality was recorded at 48 h. All final counts were made with a wide-bore pipette under a dissecting microscope (15x magnification) after first lowering the test solution level in the chambers.

3.2.3.3 Sheepshead Minnow

A 48-h cadmium chloride reference toxicant test was conducted

at the following nominal cadmium concentrations: 0, 0.32, 0.56, 1.0, 1.8, 3.2, 5.6, and 10.0 mg/L. Ten fish larvae (< 24-h old) were placed in replicate 600 ml glass beakers containing 400 ml of test solution. The larvae were transferred to each beaker with a wide-bore, fire-polished glass pipette. Test beakers were placed in a biological incubator to maintain constant temperature (25 C) and photoperiod (12-h L:12-h D). Two reference toxicant tests were conducted with sheepshead minnow larvae because the source of these larvae was different. One group of larvae was obtained from established cultures in our laboratory. The other source was Aquatic Bioassay Laboratory in Baton Rouge, LA.

3.2.3.4. Fathead Minnow

A 48-h cadmium chloride reference toxicity test was conducted at the following nominal cadmium concentrations: 0, 0.032, 0.056, 0.1, 0.18, 0.56 and 1.0 mg/L. Ten fish larvae (< 24-h old) were placed in replicate 600 ml glass beakers containing 400 ml of test solution. The larvae were transferred to each beaker with a wide-bore glass pipette. Test beakers were placed in a biological incubator to maintain constant temperature (25 C) and photoperiod 16-h L: 8-h D.

3.2.3.5 <u>Ceriodaphnia</u>

A 48-h cadmium chloride reference toxicity test was conducted at the following nominal cadmium concentrations: 0, 0.018, 0.032, 0.056, 0.10, 0.180, 0.320 and 0.560 mg/L. Five Ceriodaphnia

neonates (< 4-h old) were pipetted into replicate 30 mL glass beakers containing 15 mL test solution (10 neonates per condition). Neonates were counted by drawing them into a narrow-bore glass pipette and examined under a dissecting microscope (15x magnification). Test beakers were placed in a biological incubator to maintain constant temperature (25 C) and photoperiod 16-h L: 8-h D. The organisms in each test beaker received 0.1 mL Selenastrum and 0.24 mL CEROPHYL initially and at 24 h.

3.2.4 Sample Collection, Handling and Storage

Ambient water was collected from all eight stations and transported to a toxicity testing facility at the Wye Research and Education Center. Grab samples collected from each station were a composite of the water column (~ 1m from the top and ~ 1m from the bottom). A metering pump (12 V DC Little Giant Utility Pump, Model PPS-12) with polyethylene line was used to collect samples in 11.25 L glass containers.

The time lapsed from the collection of a grab sample and the initiation of the test or renewal did not exceed 72 h. Samples were collected on day 0, day 3 and day 6 during the 8-day tests. All samples were chilled after collection and maintained at 4 c until used. The temperature of the ambient water used for testing was 25 C. Salinity adjustments (increase) were performed on samples collected from saline sites to obtain a standard test salinity of 15 ppt. Slightly higher ambient salinity conditions of 19-20 ppt were reported at the Elizabeth River station;

therefore, salinity was not changed.

3.2.5 <u>Test Procedures</u>

3.2.5.1 Grass Shrimp

Eight to 9-d old larvae were exposed to each ambient water condition from all eight stations and a 15 ppt salinity control. Salinity adjusted freshwater (15 ppt) from the Indian Head, Freestone Point and Possum Point stations (Potomac River) was used. During the first set of experiments (Elizabeth River, Patapsco River and Wye River), larvae were tested individually in Nitex chambers (0.202 mm mesh) to prevent cannibalism. Each test condition consisted of 4 larvae per 600 ml beaker with 12 beakers (total of 48 larvae per condition). Each beaker contained 200 ml of sample water; no aeration was supplied to the beakers. Larvae were also tested in groups of 5 at two test conditions during these initial tests to determine if cannibalism occurred. Cannibalism did not occur, therefore, all future tests (Potomac River experiments) were conducted with groups of 5 larvae per 600 ml Eight beakers were used for each test condition for a beaker. total of 40 larvae per condition.

Temperature, salinity, dissolved oxygen and pH were measured daily in randomly-selected beakers at each test condition prior to water exchange. One beaker was randomly selected per treatment condition daily to monitor ammonia. One-half of the water (100 ml) was siphoned out daily from each beaker and then renewed with a fresh sample held at 4 C and warmed to 25 C. Larvae were fed

concentrated volumes of <u>Artemia</u> nauplii daily after water renewal. For larvae tested individually, 75 ul of concentrated <u>Artemia</u> was used daily. Approximately 375 ul of concentrated <u>Artemia</u> was fed to larvae tested in groups of five. At the completion of the 8-d experiments, all larvae were preserved in 10% formalin prior to wet weight analyses. Dry weights for each larvae were recorded following 72-h drying at 70 C.

Survival proportions were compared using the Kruskal-Wallis test. A nonparametric test was chosen in lieu of ANOVA because the homogeneity of variance assumption was not satisfied in the data set. Growth parameters were compared using ANOVA and the Bonferroni T-test. This t-test was chosen because of the unequal replicate numbers (Weber et al., 1989). Differences between means were considered significant at the p< 0.05 level.

3.2.5.2 Copepod (E. affinis)

The methods used for testing <u>E</u>. <u>affinis</u> were modified from Hall et al., 1988b. Experiments were started with <u>E</u>. <u>affinis</u> nauplii <24-h old. Nauplii were exposed to all eight test conditions and an artificial seawater control (15 ppt). Salinity adjusted freshwater 15 ppt from the Indian Head, Freestone Point and Possum Point stations (Potomac River) was used. Four replicate 150 ml beakers containing 100 ml of test water were used for each condition. Ten to fifteen nauplii were placed in a chamber suspended within each test beaker. The initial number of nauplii were recorded. The chamber design was the same as that used in the

reference toxicant tests described previously (Section 3.2.3.2). Nauplii were counted using a wide-bore, glass pipette under a dissecting microscope (15x magnification). Test beakers were held in a biological incubator to control temperature (25 C) and photoperiod (12-h L:12-h D). Temperature, dissolved oxygen, and salinity were measured daily in selected beakers. On alternate days, pH values were recorded. A one liter sample was taken daily from each of the ambient test conditions, stored at 4 C, and warmed to 25 C. One-half of the water volume per beaker was siphoned out daily and renewed with the fresh sample. Following the daily water renewals, copepods were fed an equal volume mixture of $\underline{\mathrm{T}}$. fluviatilis and I. galbana. Four mL of the two-species algal mix, each in log-phase, was pipetted into each test chamber. Algal cell densities within the chambers were generally 2 - 5 \times 10⁵ cells/ml. All cell counts were done with Spencer improved Neubauer corpuscle counting chamber (hemacytometer).

On day four of exposure to ambient water, two test beakers from each condition were selected to evaluate nauplii survival. Counts were made by first lowering the volume of water in the chambers, then removing the remaining water and nauplii in small aliquots with a wide-bore glass pipette. The nauplii were counted under a dissecting microscope (15x magnification) then discarded. Survival was evaluated again on day eight using the two remaining beakers at each test condition. Surviving egg-carrying adult female copepods were isolated in 20 ml glass vials (l adult/vial) from each test condition. Within 24-48 h, egg-sacs had been

released by the females and the resulting brood size was determined for each female.

Percent survival of copepods was compared using the Kruskal-Wallis test. A nonparametric test was used in lieu of ANOVA because the homogeneity of variance assumption was not satisfied in the data set. Analysis of variance (ANOVA) and the Bonferroni T-test were used to compare brood sizes. Differences between means were considered significant at the p< 0.05 level.

3.2.5.3 Sheepshead Minnow

Experiments were initiated with sheepshead minnow larvae < 24-h old. Larvae were exposed to all test conditions and an artificial seawater control (15 ppt). Four replicates of 10 larvae per replicate (600 ml beaker) were used for the Potomac River tests. However, due to poor hatching success prior to the first set of experiments, the number of larvae per test condition was reduced. Three replicate 600 ml glass beakers, per condition, containing 400 ml of test water were used for the control and Elizabeth River conditions. Four replicate beakers per condition were used for Patapsco River and Wye River conditions. Each of the beakers contained eight larvae with the exception of one replicate from the Elizabeth River condition which had only seven larvae. Larvae were transferred to the beakers with a wide-bore, fire-polished glass pipette.

Temperature, salinity, and dissolved oxygen were measured daily in selected beakers prior to water exchange. On alternate

days, pH measurements were recorded. Test beakers were held in a biological incubator to maintain constant temperature (25 C) and photoperiod (12-h L:12-h D). A one liter sample was taken daily from each of the three field samples, stored at 4 C, and warmed to 25 C. One-half of the water volume per beaker was siphoned out daily and renewed with the fresh sample. Following the daily water renewals, larvae were fed <24-h old <u>Artemia</u> nauplii. <u>Artemia</u> nauplii were concentrated on a Nitex mesh and rinsed briefly with culture water (15 ppt artificial seawater). Four mL of the concentrate was diluted to 100 mL with culture water. One-half mL of the diluted nauplii was pipetted into each beaker.

Larval survival was recorded daily for eight days. At the end of 8-d exposure to ambient water, one-half of the surviving larvae from each beaker were preserved in 10 % formalin. The remaining larvae were rapidly frozen in liquid nitrogen for protein analysis (see Suborganismal Section). Larvae were removed from the formalin and rinsed with deionized water prior to measuring wet weight and length. Dry weights for pooled larvae samples were recorded following 31-h drying at 78 C. Mean individual larval weights were calculated from the pooled weights.

The average dry weight of control larvae must exceed 0.25 mg/L for an acceptable growth test. Our SOP's state that an average dry weight of 0.50 mg is required for an acceptable test. This value is based on culture and testing salinities of 20 - 30 °/oo. The salinity used in these tests (15 °/oo) is much lower and results in lower growth rates.

Survival proportions were compared using the Kruskal-Wallis test. A nonparametric test was used because the homogeneity of variance assumption was not satisfied in this data set. Some conditions had zero variance. Growth parameters were compared using ANOVA and the Bonferroni T-test. Differences between means were considered significant at the p<0.05 level.

3.2.5.4. Fathead Minnow

Experiments were initiated with fathead minnow larvae less than 24-h old. Larvae were exposed to the following test conditions: Control water, and Potomac River water collected from Indian Head, Freestone Point and Possum Point. Four replicate 600 ml glass beakers containing 400 mls of test water were used for each condition. Ten larvae were transferred to each beaker with a fire-polished, wide-bore glass pipette. Beakers were placed in a biological incubator to maintain constant temperature (25 C) and photoperiod (16-h L: 8-h D). The larvae in each test beaker were fed 0.1 mL of a concentrated suspension of newly hatched brine shrimp nauplii twice daily. Nauplii were rinsed with fresh water prior to feeding larvae.

Temperature, dissolved oxygen, and pH were measured daily in selected beakers from each condition. Hardness was measured three times corresponding to each new sample collected from the Potomac River. Larval mortality was recorded daily and dead larvae were discarded. A one-liter sample was taken daily from each of the three field samples, stored at 4 C, and warmed to 25 C. Before the

renewal of test solutions, uneaten and dead brine shrimp were removed from the bottom of each test beaker. The test solutions were renewed daily immediately after cleaning each beaker. One-half the water volume per beaker was siphoned out and renewed with the fresh, warmed sample.

At the end of 8-d exposure to ambient water, one-half of the surviving larvae from each beaker were preserved in 10% formalin. The remaining larvae were rapidly frozen in liquid nitrogen for subsequent protein analysis. Larvae were removed from the formalin and rinsed with deionized water prior to measuring wet weight and length. Dry weights for pooled larvae samples were recorded following 31-h drying at 78 C. Mean individual larval weight was calculated from the pooled weight.

Survival proportions were transformed using the arc sine (square root (Y)) transformation. Analysis of variance (ANOVA) and Dunnetts Test were used to compare survival in the river conditions to the control. Growth parameters were compared using ANOVA followed by Dunnetts Test or Bonferroni T-Test. Differences between means were considered significant at the p<0.05 level.

3.2.5.5 <u>Ceriodaphnia</u>

Experiments were initiated with <u>Ceriodaphnia</u> neonates less than 24-h old and released during the same 4-h period. Neonates were exposed to the following test conditions: Control water, and Potomac River water collected from Indian Head, Freestone Point and Possum Point. Ten replicate 30 mL glass beakers containing 15 mLs

of test water were used for each of the four conditions. One neonate was placed in each beaker with a narrow-bore glass pipette. Beakers were placed in a biological incubator to maintain constant temperature (25 C) and photoperiod (16-h L: 8-h D). During the test, the <u>Ceriodaphnia</u> were fed the same diet as used for the cultures. The organisms in the test beakers were fed 0.1 mL <u>Selenastrum</u> and 0.24 mL CEROPHYL daily.

Two <u>Ceriodaphnia</u> tests were conducted because the mean number of neonates produced per adult in the control during the first experiment was less than 15. A minimum number of 15 is needed for a valid test using reproduction as an endpoint. However, survival data from the first experiment was valid.

Temperature, dissolved oxygen, and pH were recorded daily in selected beakers from each test condition. Hardness was measured three times corresponding to each new sample collected from the Potomac River. A glass pipette was used to transfer each test organism daily to a new beaker containing 15 mL of fresh sample to which food had already been added. Prior to transferring each organism, adult mortality and neonate production was recorded daily. The young were counted with the aid of a dissecting microscope (15x magnification). Fishers Exact Test and/or a T-test were used to analyze survival (Weber et al., 1989). Analysis of variance (ANOVA) and the Bonferroni T-Test were used to compare reproduction. Differences between means were considered significant at the p<0.05 level.

3.2.5.6 Fundulus heteroclitus

F. heteroclitus exposed to ambient water from all saline sites were used for suborganismal testing (protein analysis). These fish were collected near the Chesapeake Biological Laboratory and transported to our laboratory by Dr. Jay Gooch. Test organisms were exposed to all 5 ambient saline conditions and an artificial seawater control (15°/00) at 25 C. Replicate 18 L glass aquaria with 14 L of water contained six fish per aquaria. One-half of the water volume in each test aquaria was siphoned out daily and renewed with fresh sample. Aeration was supplied if dissolved oxygen fell below 5.0 mg/L. Temperature, salinity and dissolved oxygen were measured daily and on alternate days pH values were recorded. Fish were fed once daily to satiation with flake food (Tetramin®). Experiments were terminated after 8 days and surviving fish were given to Dr. Gooch for protein analysis.

3.2.6 Contaminant and Water Quality Evaluations

The contaminant analyses used for these studies provided limited information on selected contaminants present in the study areas. It was not our intention to suggest that the proposed analysis for inorganic and organic contaminants will provide an absolute cause and effect relationship between contaminants and biological effects if effects were reported. Information on suspected contaminants in the study areas would, however, provide valuable insights if high potentially toxic concentrations of contaminants were reported in conjunction with biological effects.

Aqueous samples for analysis of organic and inorganic contaminants listed in Table 2 were collected during the ambient toxicity tests. Analytical procedures and references for analysis of these samples are presented in Table 3. Total inorganic contaminant analysis was conducted on filtered samples using 0.40 um polycarbonate filters.

Four liter whole water samples were collected for organic contaminants analysis (Table 2). Organic contaminants other than those identified in Table 2 (non-target organics) were measured if GC/MS peaks were identified. Detailed procedures for preparing samples for inorganic and organic analysis are described in detail in Hall et al. (1988a). These contaminants have been evaluated during our previous striped bass contaminant studies on the Potomac River. Various metals (cadmium, lead and zinc) and organics (chlordane) have been reported at potentially toxic concentrations in the Potomac River (Hall et al., 1988a). Analysis for both organic and inorganic contaminants was conducted at least one time on aqueous samples collected from each station. If toxicity was reported during the experiments then the suite of contaminants was analyzed on at least 2 separate samples taken during these tests. VERSAR Inc. was responsible for all organic and inorganic analyses.

Standard water quality conditions of temperature, salinity, dissolved oxygen, pH, conductivity and hardness (freshwater stations) were evaluated at each site after sample collection. These conditions were evaluated every 24 h at all test conditions during the tests. Datasonde units (continuous water quality

Table 2. Concentrations of the following organic and inorganic contaminants were evaluated.

Contaminant	Detection Limit (ug/L)
Aroclor 1248	0.050
Aroclor 1254	0.050
Aroclor 1260	0.050
DDE	0.02
Toxaphene	0.2
Chlordane	0.02
Perylene	0.70
Fluorene	0.90
Phenanthene	0.70
Anthracene	0.70
Fluoranthrene	1.1
Pyrene	1.0
Benz(a)anthracene	1.7
Chrysene	0.7
Aluminum	3.0
Arsenic	3.0
Cadmium	2.0
Chromium, total	3.0
Copper	2.0
Lead	2.0
Mercury	0.2
Nickel	5.0
Selenium	3.0
Tin	5.0
Zinc	5.0

Table 3. Analytical methods used for organic and inorganic analysis. The following abbreviations were used: GC-EC (Gas Chromatography - Electron Capture), GC -MS (Gas Chromatography - Mass Spectrometry), Atomic Emission - ICP (AE-ICP), AA-H (Atomic Absorption - Hydride), AA-F (Atomic Absorption - Furnace) and AA-DA (Atomic Absorption - Direct Aspiration) and AA-CV (Atomic Absorption - Cold Vapor).

Contaminant	Method	Method #	Reference
Halogenated Hydro- carbon Pesticides	GC-EC	608	U. S. EPA, 1984
Polychlorinated Biphenyls	GC-EC	608	U. S. EPA, 1984
Base-Neutral Extractable Organic Compounds	GC-MS	625	U. S. EPA, 1984
Aluminum	AE-ICP	200.7	U. S. EPA, 1984
Arsenic	AA-H	206.3	U. S. EPA, 1979
Cadmium	AA-F	213.2	U. S. EPA,
Chromium, Total	AA-F	218.2	1979 U. S. EPA
Copper	AA-F	220.2	1979 U. S. EPA
Lead	AA-F	239.2	1979 U. S. EPA
Mercury	AA-CV	245.1	1979 U. S. EPA
Nickel	AA-F	249.2	1979 U. S. EPA,
Selenium	AA-H	270.3	1979 U. S. EPA,
Tin	AA-F	282.2	1979 U. S. EPA,
Zinc	AA-DA	289.1	1979 U. S. EPA, 1979

monitors) were used to continuously record the above water quality conditions at the various stations during the Potomac River experiments. (Appendix B)

3.3 <u>Sediment Tests</u>

3.3.1 <u>General Description</u>

All tests and analyses were conducted according to the SOPs and QA plans previously submitted to the sponsor. The methods described in this report are general summaries of those protocols. Sediment samples (100% ambient sediment samples) from eight stations were tested using 3 organisms: the grass shrimp, Palaemonetes pugio, the amphipod, Lepidactylus dytiscus, and the polychaete worm, Streblospio benedicti. All tests were conducted for 10 d at 25C and monitored daily. At the end of 10 d, mortalities were recorded, and the animals were returned to the original test containers. The organisms were then monitored daily for an additional 10 d. Numbers of live animals were recorded on day 20, and any living organisms were preserved for length/weight measurements. The sediment samples were collected from the Elizabeth River, Patapsco River, Potomac River, and the Wye River. Three of the Potomac sites were located in freshwater areas. other sites in the Potomac River, the Wye River and Patapsco River had low salinities (between 7 and 9 ppt). All of these samples were adjusted to 15 ppt by sieving with 15 ppt control water prior to testing. The eighth station, in the Elizabeth River, was 20 ppt without adjustment, and was tested at 20 ppt. Control sediments for each species consisted of the native sediments from the area in which the test organisms were collected. A control was tested with each set of samples. In addition, a reference sediment (see below) was tested with each set of test samples.

Sediment was collected for particle size analysis from each of the test stations several weeks prior to the toxicity tests. reference sediment was then chosen for each set of samples. From the initial sediment collection, the Wye River sample matched the sediment particle size needed for a reference sediment for the first set of test samples. The Patapsco River site initially sampled was less than 1% sand (99% silt/clay) and the Elizabeth River site was 5% sand (95% silt/clay). The Wye River sample was about 4.5% sand, and therefore, was suitable as a reference sediment to be used for these two sites (Table 4). However, when actual test sediments were collected, the size/composition of the Patapsco River site was drastically different from the initial sample. The test sediment was approximately 94% sand (Table 5). However, the amphipod control sediment tested was also a very sandy sediment (approx. 99% sand), so an additional reference sample was not needed. The second set of test samples fit into two categories: greater than 85% sand and greater than 85% silt/clay. A reference sediment from the Nansemond River, Virginia, was chosen for the greater than 85% silt/clay samples, and the amphipod control sediment was used for comparison to the sandy samples.

Table 4. Particle size analysis of sediments collected for pilot survey from the eight stations. The five Potomac River sites are Indian Head, Freestone Point, Possum Point, Morgantown, and Dahlgren.

Station	% Sand	% Silt	% Clay
Set #1:	3		
Sec #1:			
Elizabeth River (Atlantic Wood)	5.14	68.03	26.84
Patapsco River	0.16	92.71	7.13
Wye River	4.47	86.86	8.67
Set #2:			
Indian Head	14.43	69.23	16.34
Freestone Point	10.19	78.58	11.23
Possum Point	3.90	74.95	21.15
Morgantown	95.98	1.40	2.62
Dahlgren	89.82	4.93	5.25

Table 5. Particle size analysis of sediments from eight stations and controls used in toxicity tests. The five Potomac River sites are Indian Head, Freestone Point, Possum Point, Morgantown, and Dahlgren.

Station	<pre>% Sand</pre>	<pre>% Silt</pre>	<pre>% Clay</pre>
Set #1:			
Elizabeth River (Atlantic Wood)	13.84	37.24	48.93
Patapsco River	94.32	2.84	2.84
Wye River	20.20	48.85	30.95
Worm Control	84.05	9.50	6.45
Amphipod Control	100.00	0.00	0.00
Grass Shrimp Control	100.00	0.00	0.00
Set #2:			
Indian Head	11.18	56.85	31.97
Freestone Point	17.08	60.48	22.44
Possum Point	1.22	69.34	29.44
Morgantown	92.03	2.87	5.10
Dahlgren	85.43	7.73	6.85
Nansemond Reference	19.80	42.80	37.40
Amphipod Control	100.00	0.00	0.00

3.3.2 <u>Collection, Handling, Maintenance and Culture of Test</u> <u>Species</u>

3.3.2.1 Grass shrimp - Palaemonetes pugio

Grass shrimp were collected from the Lynnhaven Inlet, Virginia, quarantined, and identified with identifications of the species verified by a local crustacean specialist. Gravid adult females were then selected and moved to breeding tanks. As larvae were released, the adults were removed to prevent predation. Larvae were reared in 10 gallon tanks to the post-larval stage. Grass shrimp post-larvae approximately 30-d old were used in the first set of tests. Half of the animals were acclimated to 20 ppt salinity and half acclimated to 15 ppt prior to the tests.

Grass shrimp were purchased from a supplier (Multi Aquaculture Systems, New York) for the second set of tests. The shrimp were handled in the same fashion as mentioned above. We received post larvae 9-12 d old on September 25, 1990. These animals were held at 25 C and 15 ppt salinity, fed 24 hr old artemia nauplii daily ad libitum, and acclimated to laboratory conditions. The shrimp were held for observation for 4-d prior to initiation of the tests. Animals 12 to 15-d old were used in the second set of tests. Less than 10% mortality was observed during the holding and acclimation period.

3.3.2.2. Amphipod - Lepidactylus dytiscus

Animals were collected from an estuarine site at Virginia Beach, Virginia, transported in buckets to the AMRL and acclimated

to laboratory conditions. Amphipods were collected at 20 ppt salinity. Half of the population were acclimated to 15 ppt over a 4-d period, and the remaining half were maintained at 20 ppt. Animals were held in their native sediment (sand) in 20 gallon tanks for approximately 7 d prior to initiation of the tests. Less than 10% mortality was observed during the holding period. For the second set of tests, the animals were again collected from the field and brought to the laboratory approximately one week prior to initiation of the toxicity tests. These animals were acclimated to 15 ppt over a 4-d period. Again, mortality during the holding period was less than 10%.

3.3.2.3 Polychaete worm - Streblospio benedicti

Worms were collected from a tidal creek entering the Lafayette River (Norfolk, VA), brought to the laboratory, and held in 10 gallon tanks of native mud with 15 ppt overlying water approximately one week before testing. A subset of animals was acclimated to 20 ppt over a 4-d period for the first set of tests. Animals were sieved out of the holding tank and placed into 4" culture dishes 24 h prior to addition to the test containers. Both sets of organisms had less than 10% mortality during the holding period.

3.3.3 Reference Toxicant Tests

The relative sensitivities of each set of test organisms was evaluated by a reference toxicant test designed to assess changes

in tolerances of the organisms that may have been due to disease, or stress developed from handling and acclimation. The grass shrimp, Palaemonetes pugio, was tested in a 96-h, water only, acute test using SDS (sodium dodecyl sulfate) in 1 liter containers with 800 mL of test solution. SDS was selected as the reference toxicant to be used with the grass shrimp because it is a standard EPA reference toxicant with a large historical data base. Literature exists for grass shrimp tested with SDS, and there is minimal human health risk from this toxicant. Ten animals were placed into each test container, with two replicates per concentration. Each batch of organisms used for each set of bioassays was tested with a reference toxicant.

The amphipod, <u>Lepidactylus dytiscus</u>, was tested in a 96-h, water only, cadmium chloride reference toxicant test using 400 mL of solution with 10 animals per container. Five concentrations and a control (seawater only) were prepared by serial dilution. Tests were conducted at 20 ppt salinity with two replicates for each concentration.

The polychaete worm, <u>Streblospio benedicti</u>, was tested in a 96-h, water only, cadmium chloride (CdCl₂) reference toxicant test. Animals were tested in 250 mL crystallizing dishes with 200 mL of test solution, 10 animals per dish and two replicates per concentration.

3.3.4 Sample Collection, Handling and Storage

Sediment samples were collected at each site by AMRL personnel

and returned to the laboratory for testing. The first set of sediments was collected at the following sites: Elizabeth River (Atlantic Wood), Patapsco River (Bethlehem Steel), and the Wye River. All samples were collected on Thursday, August 16, 1990, by petite ponar grab. Grab samples were collected and emptied into a large stainless steel pan. The top 2 cm of sediment was then removed from each grab and placed into a tub. Enough grabs were taken at each site to fill a tub with approximately 8 gallons of sediment. Each composite sediment sample was then homogenized by stirring (using an analytically cleaned stainless steel paddle). Aliquots were taken from the tub and distributed to individual sample containers for toxicity tests, chemical analysis and particle size analysis. All samples were transported to the AMRL in coolers, on ice, and out of direct sunlight. Samples were logged into AMRL refrigerators and held at 4 C until toxicity tests were initiated. All sediment samples used for bioassays were tested within the recommended two week holding period (USEPA/ACOE, 1977). Samples for chemical analysis were frozen until ready for testing.

The second set of sediment samples was collected on September 26, 1990, at the following Potomac River sites: Dahlgren, Indian Head, Possum Point, Morgantown and Freestone Point. A sample was also collected at a Nansemond River reference site in Suffolk, VA. All samples were collected and handled the same as the first set. Approximately 8 gallons of sediments were collected by petite ponar grab, homogenized and distributed to sample containers. All

samples were transported on ice, out of direct sunlight. Samples were held in refrigerators at 4 C until used in toxicity tests. Samples for chemical analysis were stored frozen until tested. All samples were analyzed within EPA recommended holding times.

3.3.5 Test Procedures

Sediments were removed from the refrigerator and homogenized by stirring. Large sticks or debris and macrofauna were removed by sieving the sediments through a 0.5 mm mesh screen prior to placing them into test containers. A two centimeter layer of sediment was placed into each test dish with overlying water made from artificial sea salts and deionized water. The test containers were placed into water baths and allowed to equilibrate 24-h before animals were added. The first set of test sediment was placed into the test containers on August 27, 1990. All test animals were added the next day (August 28, 1990). Seven replicates were set up for each test sediment and a control. A subset of each test population of organisms was set aside for initial growth measurements. All containers were gently aerated throughout the test (no more than 100 bubbles/minute). All tests were conducted The test was monitored daily for 10 d for oxygen, temperature, pH, mortality and behavior (i.e., amphipod emergence from sediment). At day ten (September 7, 1990), the animals were counted, mortality assessed, overlying water changed, and living animals returned to test containers for an additional 10 d. Organisms from replicates 6 and 7 were removed and given to Dr. Jay Gooch, CBL, for stress protein analysis (see Section 3.4). On day 20 of the test (September 17, 1990) mortality was recorded and surviving organisms were measured for growth. The second set of sediment samples (Potomac sediment) was treated the same as the first set. The test material was placed into the test containers on September 27, 1990, and allowed to equilibrate overnight. Test organisms were added on September 28, 1990, and monitored for 10 d. On October 8, 1990, day 10 of the test, mortality was assessed and animals from replicates 6 and 7 were collected and processed for stress protein analysis. All surviving organisms in replicates 1 through 5 were returned to the test containers for an additional 10-d. On October 18, 1990 (day 20), the test was terminated, mortality recorded and growth measurements made on surviving organisms.

In addition to the three species used for toxicity tests (i.e., the shrimp, amphipod and worm) populations of mummichogs, Fundulus heteroclitis, were exposed to test solutions in five gallon aquaria with 2 cm of sediment and four gallons of overlying water. These fish were used for protein analysis by Dr. Jay Gooch of CBL. The animals were provided by Dr. Gooch and stocked six per tank. Two replicates were set up for each test sediment and the control. Tanks were monitored daily and 75% of the overlying water was changed every 48 hours for ten days. The animals were fed flake food daily. On day ten, all fish were removed and given to Dr. Gooch for analysis (see Section 3.4).

3.3.5.1 Grass Shrimp - Palaemonetes pugio

The test methods for the grass shrimp, Palaemonetes pugio, followed the methods described in the EPA/ACOE Implementation manual (1977) for sediment bioassays. Twenty juvenile grass shrimp were placed into a one gallon glass aquarium containing a 2 cm layer of sediment and 3000 mL of overlying water. All animals were fed live artemia twice a day, and 75% of the overlying water was changed every 48 h. The shrimp control sediment was clean sand from their native habitat and was tested at 20 ppt salinity. Oxygen, temperature, pH and mortality were recorded daily for 10 d. Animals were removed from the tanks and survival recorded on day ten. Replicate 6 and 7 animals were given to Dr. Gooch for protein analysis. The remaining five replicates of animals were returned to the test chambers and monitored for an additional 10 d. On day twenty, all animals were measured for length and dry weight.

3.3.5.2 Amphipod - Lepidactylus dytiscus

A series of test containers was set up according to the methods outlined in the ASTM "Guide for Conducting Solid Phase 10-Day Static Sediment Toxicity Tests with Marine and Estuarine Amphipods" (ASTM, 1990). Two centimeters of sediment were placed into each 1 liter jar with 700 mL of overlying water. Twenty animals were placed into each test jar and monitored for 10 d. Control sediment was native sand from the amphipod collection site and was tested at 20 ppt salinity. A subset of the population of

test animals was selected for initial length/weight measurements. Test containers were monitored daily for oxygen, temperature, and ph. Observations relative to the number of animals emerged from the sediment were also recorded. The amphipods were not fed during the test. At the end of 10 d, animals were sieved from test containers, mortality recorded, and surviving animals returned to the test containers for an additional 10-d period. On day twenty, animals were again sieved from the containers and mortality recorded. Live animals were preserved for growth measurements.

3.3.5.3 Polychaete worm - Streblospio benedicti

Test methods for the polychaete, <u>Streblospio benedicti</u> are similar to those applied to the amphipod test. Ten organisms were placed into each 750 mL test jar containing 2 cm of sediment and 500 mL of overlying water. Test containers were monitored daily for oxygen, temperature and pH. The worms were not fed during the test. After 10 d, the worms were sieved from the sediment and mortality recorded. Surviving organisms were returned to the test dishes and monitored for an additional 10 d. On day twenty, the worms were sieved from the test jars, counted and preserved for growth measurements.

3.3.6 Contaminant and Sediment Quality Evaluations

3.3.6.1 Semi-Volatile Organic Analysis: Sediment

The sediment sample and corresponding QA/QC spikes and blank were analyzed for semi-volatile (base/neutral and acid

extractables) organic compounds (Table 6) in accordance with USEPA methods 3550 and 8270 (USEPA, 1986). In summary, the method requires an ultra-sonication extraction of approximately thirty grams of sample with a methylene chloride:acetone solvent system. Each sample was sonicated three times and the filtrates pooled as the final extract. Each extract was then concentrated using Kuderna-Danish and rotary evaporation techniques to a final volume of 1 ml. Each sample received six USEPA-approved surrogates and, after concentration, each 1 ml extract was spiked with six USEPA-approved internal standards.

Sample extracts were analyzed using a Finnigan INCOS-50 GC/MS/DS. The MS was tuned to USEPA criteria by injecting 50 ng of decafluorotriphenylphosphine (DFTPP) into the GC inlet for tune criteria.

A calibration curve for all analytes was established using a five-point calibration. All analytes were calibrated at 12.5, 25, 50, 100, and 200 ng/uL using the internal standard method. The calibration curve was checked and corrected regularly using the 100 ng calibration point.

Tentative identifications were made based on the USEPA accepted National Bureau of Standards (NBS) compound library. The substances of greatest apparent concentration (to a maximum of 20) are reported for each semi-volatile organic sample.

The GC was equipped with a J&W DB-5 (30m x .32mm ID x .25 μ m film) capillary column and sample introduction was via a grab type split/splitless injection system. The MS scanned from 35 to 450

Table 6. Semi-volatile organic compounds analyzed for utilizing a user-created calibration library. Sediment method detection limits (MDL) are reported in $\mu g/kg$ dry weight.

CAS NO.	COMPOUND	SEDIMENT	MDL
65-53-3	Aniline	14.5	
95-57-8	2-chlorophenol	13.2	
111-44-4	Bis(2-chloroethyl)ether	11.2	
108-95-2	Phenol	11.9	
541-73-1	1,3-dichlorobenzene	11.9	
106-46-7	1,4-dichlorobenzene	12.5	
95-50-1	1,2-dichlorobenzene	11.9	
100-51-6	Benzyl alcohol	27.1	
39638-32-9	Bis(2-chloroisopropyl)ether	5.9	
95-48-7	2-methylphenol	15.8	
91-57-6	2-methylnaphthalene	9.2	
67-72-1	Hexachloroethane	21.8	
621-64-7	n-nitroso-di-n-propylamine	13.2	
106-44-5	4-methylphenol	13.9	
98-95-3	Nitrobenzene	11.2	
78-59-1	Isophorone	6.6	
88-75-7	2-nitrophenol	27.1	
65-85-0	Benzoic acid	18.5	
105-67-9	2,4-dimethylphenol	15.8	
111-91-1	Bis(2-chloroethoxy) methane	9.9	
120-83-2	2,4-dichlorophenol	21.8	
120-82-1	1,2,4-trichlorobenzene	15.2	
91-20-3	Naphthalene	4.6	
106-47-8	4-chloroaniline	26.4	
87-68-3	Hexachlorobutadiene	22.4	
59-50-7	4-chloro-3-methylphenol	20.5	;
77-47-4	Hexachlorocyclopentadiene	25.7	
88-06-2	2,4,6-trichlorophenol	37.0)
95-95-4	2,4,5-trichlorophenol	44.9)
88-74-4	2-nitroaniline	37.6	;
91-58-7	2-chloronaphthalene	9.9	
208-96-8	Acenaphthalene	5.9	
84-66-2	Dimethylphthalate	9.9	
606-20-2	2,6-dinitrotoluene	48.2	?
99-09-2	3-nitroaniline	247	
83-32-9	Acenaphthene	9.9	
51-28-5	2,4-dinitrophenol	262	
132-64-5	Dibenzofuran	7.9	
100-02-7	4-nitrophenol	268	
121-14-2	2,4-dinitrophenol	43.6	5
86-73-7	Fluorene	9.9	
7005-72-3	4-chlorophenylphenylether	20.5	5
84-66-2	Diethylphthalate	9.9	
100-01-6	4-nitroaniline	279	
534-52-1	4,6,-dinitro-2-methylpheno	1 122	
86-30-6	n-nitrosodiphenylamine	19.1	L

Table 6. (Continued)

CAS NO.	COMPOUND	SEDIMENT MDL
101-55-3	4-bromophenylphenylether	41.6
85-01-8	Phenanthrene	9.2
118-74-1	Hexachlorobenzene	37.6
87-86-5	Pentachlorophenol	136
120-12-7	Anthracene	9.9
84-74-2	Di-n-butylphthalate	5.9
206-44-0	Fluoranthene	10.6
129-00-0	Pyrene	10.6
85-68-7	Butylbenzylphthalate	17.8
56-55-3	Benzo(a)anthracene	17.8
218-01-9	Chrysene	14.5
91-94-1	3,3'-dichlorobenzidine	101
117-81-7	Bis(2-ethylhexyl)phthalate	12.5
117-84-0	Di-n-octylphthalate	7.3
205-99-2	Benzo(b) fluoranthene	13.9
207-08-9	Benzo(k) fluoranthene	13.9
50-32-8	Benzo(a)pyrene	15.2
193-39-5	Indeno(1,2,3-cd)pyrene	16.5
53-70-3	Dibenz(a,h)anthracene	17.8
191-24-2	Benzo(ghi)perylene	16.5
103-33-3	Azobenzene	7.3
92-87-5	Benzidine	24.4

amu. The GC was temperature programmed, starting at 35C then to 300C at 7C/min and held until all compounds eluted. The detection limits for calibrated priority pollutants are presented in Table 6. The detection limit for tentatively identified non-priority pollutants was determined to be 10 ug/L.

3.3.6.2 Pesticide Analysis: Sediment

The sediment sample was analyzed for organochlorine pesticides (OCP) as well as polychlorinated biphenyls (PCBs) in accordance with USEPA Methods 3550 and 8080 (Table 7). In summary, the method requires an ultra-sonication extraction of approximately thirty grams of sample with a methylene chloride: acetone solvent system. Each sample was sonicated three times and the filtrates pooled as the final extract. Each extract was then exchanged to hexane and concentrated using Kuderna-Danish and rotary evaporation techniques to a final volume of 1 mL. The extracts were fractionated using florisil column chromatography and treated with activated copper to reduce sulfur contamination. One microliter of each extract was pesticides and PCBs by organochlorine analyzed for chromatography/electron capture (GC/ECD) detection. The GC was equipped with a DB-17 capillary column (.32mm x 30m x .25 μ m film), and analyses were carried out isothermally at 220° C. Quantitation was based on the external standard technique (r>0.995 for all analytes of interest).

3.3.6.3 Acid Volatile Sulfides

Table 7. Method detection limits for organochlorine pesticides and PCBs. Detection limits for sediment are reported in $\mu g/kg$ dry weight.

CAS NO.	COMPOUND	SEDIMENT MDL
391-84-6	α-BHC	0.714
3 01- 85 - 7	β-BHC	0.559
391-86-8	δ-BHC	1.062
58-89-9	Lindane	0.616
76-44-8	Heptachlor	0.819
309-00-2	Aldrin	0.608
1024-57-3	Heptachlor epoxide	0.570
959-98-8	Endosulfan I	0.859
60-57-1	Dieldrin	0.898
72-55-9	4,4'-DDE	0.528
33213-65-9	Endosulfan II	0.745
72-20-8	Endrin	1.240
72-54-8	4,4'-DDD	0.469
1031-07-8	Endosulfan sulfate	1.500
50-29-3	4,4'-DDT	3.420
72-43-5	Methoxychlor	5.0
57-74-5	Chlordane	5.0
80001-35-2	Toxaphene	10.0
2385-85-5	Mirex	1.000
7421-93-4	Endrin aldehyde	2.410
12574-11-2	Aroclor 1016	16.6
11104-28-2	Aroclor 1221	16.6
11141-16-5	Aroclor 1232	16.6
53469-21-9	Aroclor 1242	16.6
12672-29-6	Aroclor 1248	16.6
11097-69-1	Aroclor 1254	16.6
11096-82-5	Aroclor 1260	16.6

Sediment samples were frozen until analysis for acid volatile sulfides (AVS). Samples were thawed, then re-homogenized by gently stirring. Sediment samples were analyzed for AVS using the method of DiToro et al. (1990). Samples were analyzed at least in duplicate.

The analytical apparatus was a one-way flow-through system which consisted of a nitrogen purifier (deoxygenator) followed by a series of four flasks connected by glass and tygon tubing. The first flask for placement of the sample or standard was followed by a flask containing 175-200 mL of 0.05M potassium hydrogen phthalate. The last two flasks in the series each contained 175-200 mL of 0.1M silver nitrate.

Prior to analysis, the system was deoxygenated by purging with nitrogen for one hour. After purging, 10-15 grams (by weight) of sediment samples were weighed then placed into the first flask. The sample was quantitatively transferred using deoxygenated purified water. The system was then purged with nitrogen for an additional ten minutes. The AVS extraction was initiated by the addition of 6M hydrochloric acid to the sample to achieve a final hydrochloric acid molarity of 0.5M. The system was then purged with nitrogen for one hour. Sample flasks were swirled every five to ten minutes throughout the extraction. After one hour, the flow of nitrogen The silver sulfide precipitate in the third flask was stopped. (first flask originally containing silver nitrate) was filtered onto a Whatman GF/C glass fiber filter which was pre-rinsed, dried at 104C, cooled, and pre-weighed. After the silver sulfide precipitate was concentrated onto the filter, the filter was dried at 104C, cooled then re-weighed. Sample blanks were analyzed in the same manner as samples except acid was not added to the sample flask.

Sediment dry weight: wet weight ratios were calculated using separate aliquots of each sample. AVS was calculated on a micromolar basis, based upon uM of silver sulfide residue and normalized to a sediment dry weight basis.

3.3.6.4 Total Organic Carbon

Sediment subsamples were dried at approximately 50C for Total Organic Carbon (TOC) analysis. Sample homogeneity was achieved by grinding with an acid-cleaned mortar and pestle. After drying and homogenization, approximately 5 mg subsamples were placed onto precleaned sample cups and weighed using a micro-balance. Inorganic carbon was removed using 0.5M hydrochloric acid. Samples were analyzed by high temperature combustion using a Carlo-Erba NA1500 isothermal gas chromatograph in accordance with the manufacturer's instructions.

3.3.6.5 Ammonia and Nitrite

Pore water samples were removed from the sediment samples by squeezing with a nitrogen press. Water samples were filtered then frozen until analysis for ammonia-N and nitrite-N. Samples were analyzed for ammonia-N using the automated phenate method (SIC, 1981). Nitrite-N analyses were preformed using the Diazotization

Method (APHA, 1985).

3.3.6.6. <u>Metals</u>

All sediment samples were analyzed for Al, Cd, Cr, Cu, Pb, Ni, Sn, and Zn using an ICP following USEPA/SW-846, Method 6010. The elements were extracted from a thoroughly homogenized sample using 1:1 redistilled nitric acid, 30% peroxide, and concentrated hydrochloric acid as prescribed in USEPA/SW-846, Method 3050. An aliquot of the digestate was further prepared for the analysis of As and Se using a continuous hydride generation system as described in Standard Methods for the Determination of Water and Wastewater, Method 3114C. Mercury was analyzed using the cold vapor technique as described in USEPA 245.5, modified for the Contract Laboratory Program.

3.4 <u>Suborganismal Tests</u>

3.4.1 <u>Fundulus-Water and Sediment Exposures</u>

Fundulus heteroclitus used in water and sediment exposures were collected from St. John's Creek, a tidal tributary of the Patuxent River approximately one mile from the Chesapeake Biological Laboratory (CBL). Fish were collected using baited minnow traps and held in flowing, filtered, ambient Patuxent River water in tanks at CBL. In order to remove any potential influence of aromatic hydrocarbon exposure from the collection site on monooxygenase activity (unknown, though not believed to be substantial), fish were held for a minimum of 2-3 months before

use.

Detailed studies by Kloepper-Sams and Stegeman (1989) have shown that induced EROD activity in <u>Fundulus</u> readily declines following the removal of a model PAH inducer. Numerous other studies with fish species have shown this as well (for review see Payne et al., 1987). Thus, though we did not measure EROD activity in <u>Fundulus</u> prior to the start of tests, we feel confident that the holding time was sufficient to remove the influence of any potential prior exposure in St. John's Creek. For year two studies, we will test this hypothesis by evaluating EROD activity in freshly collected, versus laboratory held, <u>Fundulus</u> from this location.

At the initiation of water and sediment exposures, fish from the laboratory stock were transported in aerated containers to the laboratories at WREC and ODU, respectively. Methods for exposing Fundulus to water (Section 3.2.5.6) and sediment (Section 3.3.5) were previously described.

3.4.2 Collection of Feral Fishes

Various species of fish were collected from the sampling locations using a combination of seining, minnow traps, hook and line and trawling. We had originally intended to collect <u>Fundulus</u> <u>heteroclitus</u> from all estuarine sampling stations using minnow traps and seining, but were unable to do so, primarily due to unsuitable habitat proximate to some of the locations (i.e., the Patapsco River and the Morgantown and Dahlgren stations). At

stations where <u>Fundulus</u> were not available, and at the freshwater stations in the Potomac, we collected spot (<u>Leiostomus xanthurus</u>) and/or white perch (<u>Morone americana</u>) using a combination of hook and line and trawling with a 16 foot otter trawl. A complete list of all fish collected is contained within Appendix A. We collected approximately 125 fish of various species, from which liver samples were taken for cytochrome P-450 dependent monooxygenase activity analyses.

As fish were collected, they were brought on-board the boat, dissected, and livers from individual fish were removed and immediately frozen in liquid nitrogen. Previous studies have shown that these storage conditions are adequate to preserve monooxygenase activity for greater than one year (J. Stegeman, personal communication). The remainder of the fish were packaged in whirl-pak bags and transported to the laboratory on ice for measurements of body size (wt), sex determination and evaluation of gonadosomatic indices (gonad wt./body weight).

3.4.3 Sampling of Organisms from Water and Sediment Toxicity Tests 3.4.3.1 Fundulus

On the day toxicity tests were being terminated, <u>Fundulus</u> from the replicate test chambers were pooled, tissues dissected and frozen in liquid nitrogen as described for the feral fish samples. In general, samples from individual fish were kept separate at this point in order to be able to generate individual hepato-somatic and gonado-somatic indices. However, due to limitations in our ability

to process tissue samples with weights below approximately 200 mg, samples within treatments were generally pooled for the biochemical preparations. We processed approximately 67 samples (including pooled samples) from these tests.

3.4.3.2 Larval Fish and Invertebrates

Samples of surviving larval fish and invertebrates from water column tests were sampled from individual test chambers, pooled and frozen and stored in liquid nitrogen. We obtained samples of larval sheepshead and fathead minnows as well as grass shrimp and <u>E</u>. affinis. These samples are currently being processed.

3.4.3.3 Biochemical Methods

Microsomes were prepared for monooxygenase system measurements from tissue samples using methods described by Stegeman et al. (1987). Ethoxyresorufin-0-deethylase assays were conducted using the fluorometric technique described by Burke and Mayer (1974). This enzymatic activity is catalyzed by the cytochrome P450IA isoform in fishes (Stegeman, 1989) which is induced by planar aromatic hydrocarbon exposure. EROD activity has been widely used in field studies of fish responses to aromatic contaminants in ambient settings (Kleinow et al., 1987; Jimenez and Stegeman, 1990) and its induction is widely accepted as an indication of biologically significant aromatic hydrocarbon exposure.

Quantitation of the cytochrome P450IA1 isozyme will be done using "Western" blots (Towbin et al. 1989) using a monoclonal

antibody (1-12-3) against the P450IA1 (cytochrome P450E) of scup, a marine teleost. Kloepper-Sams et al. (1987) and Stegeman (1989) have shown that this antibody is highly specific for this P450 form in all teleosts examined to date. Relative concentrations of P450E equivalents will be determined using purified P450E from scup as a standard (Kloepper-Sams et al., 1987).

3.4.3.4 Stress Proteins

Currently, two major approaches to evaluating stress protein production (induction) exists. The most widely applied method reflects active synthesis as measured by the uptake and incorporation of a radiolabeled amino acid (e.g. [35]methionine) into protein (see for example, Bradley et al., 1988, Roccheri et al., 1981, Heikkila et al., 1982, Sanders, 1988). We quickly realized this methodology was inappropriate for our study due to cost and logistical constraints on isotope use among institutions. For example, in order to conduct isotopes based assays, each facility and P.I. would be required to be licensed for isotope use and provide the appropriate isolated work and containment areas. Alden and Mr. Hall do not currently use isotope As Dr. methodologies for their work, they do not meet these requirements. The costs and time required to do so were prohibitive relative to the time and needs of the other tests. We, therefore, did not pursue this option further and suggest that it is not a workable design for a study of this type (i.e., varying locations for tests, We also found significant problems with movement of etc.).

isotopes among locations. Finally, current guidelines for tracking isotope use and disposal would have required a personnel effort we could not meet. Since this portion of the study was only a small portion of the overall study plan and its requirements would have been disruptive to the primary tests, we chose to delete it in favor of attempting antibody based approaches which do not have these overwhelming logistical problems. We suggested that this may be a better option in the original proposal. Our current plans are to evaluate stress protein production in collected samples collected using commercially available reagent antibodies for specific stress proteins (hsp70 and hsp90).

SECTION 4

RESULTS

4.1 Water Column Tests

The following results from water column tests are presented below: toxicity data, contaminants data, water quality data and toxicity data from reference toxicant tests.

4.4.1 Toxicity Data

Survival, growth and reproduction data from the three estuarine tests conducted from 8/14/90 - 8/21/90 in the Elizabeth River, Patapsco River, Wye River stations and controls are presented in Tables 8 and 9. Survival of E. affinis was significantly lower (55%) in the Elizabeth River after 8 d when compared with the controls or other stations. However, mean brood size of the remaining E. affinis at this station was not significantly lower than the other two ambient stations or the Mean brood size from the controls was significantly controls. lower than the three ambient stations. The reason for lower reproduction in the controls may be related to the available phytoplankton and nutrients in the various conditions. water contained phytoplankton and nutrients in addition to the concentrations applied during feeding. The controls only contained phytoplankton and nutrients introduced during feeding. quantity and quality of food is important in reproduction of E.

Survival data from E. affinis, sheepshead minnow larvae and grass shrimp larvae tests conducted in the Elizabeth River, Patapsco River, Wye River and control. Tests were conducted from 8/14/90 - 8/21/90. Table 8.

Species	Station		0	m m	% Survival Per Day 4 5	. Per Day 5	v	7	ω
E. affinis	Eliz. R.				82				55*
	Pat. R.				87				92
	Wye R.				100				æ
	Control				93				100
Sheepshead	Eliz. R.	100	96	91	91	91	91	- 61	91
minnow	Pat. R.	100	100	100	100	100	100	100	100
	Wye R.	97	16	06	06	06	06	06	06
	Control	100	100	100	100	100	100	100	100
Grass	Eliz. R.	100	42	23	17	15	13	13	13*
dwraus	Pat. R.	100	86	86	64	61	61	61	61 *
	Wye R.	100	100	86	96	94	94	95	92
	Control	100	100	100	100	100	100	100	86

* Indicates value is significantly different than the control p <0.05.

* Copepods from ambient water were inadvertently introduced into test chambers.

Table 9. Reproduction (brood size) and growth data from E.

affinis, sheepshead minnow larvae, and grass shrimp
larvae tests conducted in the Elizabeth River, Patapsco
River, Wye River and control. Tests were conducted
8/14/90 - 8/21/90.

. affinis Bro	od Size Com	parisons Following 8-d E	xposures
Station	<u>n</u>	Mean Brood Size	S. E
Eliz. R.	5	64.6*	15.2
Pat. R.	10	66.1*	10.2
Wye R.	5	47.8*	7.4
Control	6	10.7	3.7
Sheepshead M	innow and G	rass Shrimp Growth Compa	risons
Sheepshead m	innow lengt	<u>h</u>	
Station	<u>n</u>	Mean (mm)	<u>S.</u> E
Eliz. R.	11	5.73	0.16
Pat. R.	16	6.13	0.22
Wye R.	14	5.61	0.23
Control	12	6.08	0.23
Sheepshead r	ninnow dry w	veight	
<u>Station</u>	<u>n</u> (poole	ed samples) <u>Mean (mg)</u>	S. I
Eliz. R.		coups of 3) 0.44	0.06
Pat. R.		coups of 4) 0.46	0.04
Wye R.		coups of 3) 0.34	0.0
Control	3 (3 g)	coups of 4) 0.30	0.1
Grass shrim	dry weight		
<u>Station</u>	<u>n</u>	Mean (mg)	<u>s.</u>
Eliz. R.	5	0.34	0.0
Pat. R.	11	0.40	0.0
Wye R.	12	0.48*	0.0
Control	12	0.41	0.0

^{*} Indicates value is significantly different than control, (p <0.05).

affinis, food in ambient conditions may account for the increased reproduction. Masters et al. (1991) have suggested a similar explanation for a freshwater zooplankton species (C. dubia) experiencing lower reproduction in control water than ambient water.

Survival, length and dry weight of sheepshead minnow larvae was not significantly reduced in ambient water from any station when compared with the controls. In contrast, survival of grass shrimp was significantly reduced at the Elizabeth River (13%) and Patapsco River (61%) stations after 8-d exposure. Survival of grass shrimp in the Wye River and control conditions was \geq 92%. Dry weight of grass shrimp was not significantly lower at any ambient station when compared with the controls. The mean dry weight at the Wye River station (0.48 mg) was, however, significantly greater than the other stations.

Survival, growth and reproduction data from estuarine and freshwater tests conducted at the five Potomac River stations and controls from 9/24/90 -10/2/90 are presented in Tables 10 - 12. There was no significant difference among survival of \underline{E} . affinis after 8 d at all stations, including controls. Number of \underline{E} . affinis produced at all five Potomac River stations was significantly greater than the controls (Table 11). The reason for lower reproduction in the controls was likely related to the quality and quantity of food as previously discussed. Survival of sheepshead minnow larvae was significantly reduced at the Morgantown and Dahlgren stations when compared to the controls.

Survival data from \underline{E} . affinis, sheepshead minnow, grass shrimp, fathead minnow and <u>Ceriodaphnia</u> tests conducted at the 5 Potomac River stations. Tests were conducted from 9/24/90-10/2/90. Salinity adjusted tests were conducted with \underline{E} . affinis and grass shrimp at the 3 freshwater stations (Possum Point, Table 10.

	Frees	Freestone Point and	Indian	Head).			1					
	Specioes	Station	7	8	Cun 3	Cumulative	₩	Survival 5	Per Day 6	7		œ
			l									
	E. affinis	Control				100						92
		Morgantown				100						96
		Dahlgren				100					•	96
		Possum Pt.*				79					ñ	100
		Freestone Pt.				81						80
		Indian Hd.				8						11
6	Sheepshead	Control	95	06	88	88		88	88	88		88
7	minnow	Morgantown	86	83	83	70		53	45	40		40*
		Dahlgren	100	95	93	06		78	09	09		20*
	Grass shrimp	Control	100	100	06	88		80	80	75		75
		Morgantown	100	95	95	93		93	93	85	,	78
		Dahlgren	100	100	100	100		86	86	86	12	*86
		Possum Pt.	100	100	100	100	1	100	86	86		*86
		Freestone Pt.	86	92	92	93		93	93	93		88
		Indian Hd.	100	86	86	86		93	93	93		88
	Fathead	Control	100	100	06	06		06	06	88		83
	minnows	Possum Pt.	100	86	06	90		06	06	06		06
		Freestone Pt.	100	86	88	88		88	88	88		88
		Indian Hd.	100	93	75	73		73	73	73		73
	+chimities increased from day	seed from day 4	to day	8 as t	this parameter		was e	evaluated	l in 2	test	chambers	ers

parameter was evaluated in 2 test ch
(see Methods Section for E. affinis) *Survival increased from day 4 to day 8 as this on day 4 and different 2 test chambers on day 8 on day 4 and different 2

Table 10 - continued

				ושניט	Cumulative &	Survival	Per Day		
Species	Station	1	8	3	4	വ	9	7	ω
Coriodaphnia	Control	100	100	100	100	100	100	68	89
5711125	Possim Pt.	100	100	100	100	06	06	80	70
	Freestone Pt.	100	100	100	100	100	100	68	78
	Indian Hd.	100	100	100	06	06	06	50	20**
1e induction	Control	100	100	100	100	100	100	100	100
Certonaphinta	Possim Pt.	100	100	100	100	100	100	100	100
a	Freestone Pt.	100	100	100	100	100	100	100	100
ξŖ	Indian Hd.	100	100	100	100	100	100	100	100
							38		
				00,00					

 1 Test conducted 10/26/90 - 11/3/90 $_{\star}$ Indicates value is significantly different than control value (p <0.05). ** Significantly different than control (p = 0.037) using a T-test.

Table 11. Reproduction of <u>E</u>. <u>affinis</u> and <u>Ceriodaphnia</u> after 8-d exposures to Potomac River water from 5 stations and the controls.

E. affinis Brood Size

Station	<u>n</u>	Mean Number of Nauplii	<u>s. e.</u>
Control	5	2.2	1.3
Morgantown	4	24*	4.4
Dahlgren	3	23.3*	5.5
Possum Pt.	8	25.4*	4.2
Freestone Pt.	6	20.8*	4.6
Indian Hd.	5	19.2*	4.0

Ceriodaphnia Neonate Production

		Mean Number of	
<u>Station</u>	n	Neonates 3 Broods	<u>S. E.</u>
Control ¹	9	1.6	0.9
Possum Pt. 1	10	5.0	0.9
Freestone Pt. 1	9	14*	1.7
Indian Hd.	10	2.0	0.7
Control ²	10	15.9	1.6
Possum Pt.2	9	16.1	1.4
Freestone Pt.2	10	17.1	1.2
Indian Hd.2	10	22.2*	1.0

^{*} Indicates value is significantly different than control value (p <0.05).

¹ Test conducted 9/24/90 - 10/2/90. ² Test conducted 10/26/90 - 11/3/90.

Table 12. Growth data for grass shrimp (dry weight), sheepshead minnow (dry weight and length and fathead minnow (dry weight and length) after 8-d exposures to Potomac River water from the 5 stations.

Grass Shrimp Dry Weight

<u>Site</u>	<u>n</u>	Mean Weight (mg)	<u>S. E.</u>
Control	8	0.59	0.03
Indian Hd.	8	0.52	0.03
Possum Pt.	8	0.53	0.02
Freestone Pt.	8	0.52	0.03
Morgantown	8	0.57	0.02
Dahlgren	8	0.58	0.02

No significant differences (p <0.05) were detected with Dunnetts Test.

Sheepshead Minnow Dry Weight and Length

<u>Site</u>	<u>n</u>	Mean Weight (mg)	<u>s. E.</u>	Length (mm)	<u>s. E.</u>
Control	9	0.25	0.02	5.8	0.3
Morgantown	8	0.29	0.10	5.9	0.4
Dahlgren	10	0.28	0.03	5.6	0.3

No significant differences (p <0.05) were detected with Bonferroni T-Test.

Fathead Minnow Dry Weight and Length

<u>Site</u>	<u>n</u>	Mean Weight (mg)	<u>s. E.</u>	Length (mm)	<u>s. E.</u>
Control	17	0.29	0.03	7.6	0.2
Possum Pt.	18	0.30	0.02	7.6	0.1
Freestone Pt.	19	0.25	0.07	7.6	0.2
Indian Hd.	15	0.36	0.04	7.8	0.1

No significant differences (p <0.05) were detected with Bonferroni T-Test.

There was no significant difference in dry weight or length among the test conditions for this species (Table 12). Survival of grass shrimp at all ambient stations was not significantly lower than the controls; survival at Possum Point and Dahlgren was significantly higher than the controls. There was no significant difference in dry weight for grass shrimp among stations.

There was no significant decrease in survival, dry weight or length of fathead minnows at the three freshwater stations when compared with the control. Survival data for Ceriodaphnia in two separate tests was different. Survival of Ceriodaphnia was significantly lower at the Indian Head Station than the control in the first experiment. In experiment 2, there was no statistical difference in survival among the 4 test conditions. The low mean number of neonates (1.6) produced in the controls for the first experiment was unacceptable for using reproduction as an endpoint; therefore, a second experiment was conducted. Reproduction data (mean number of neonates from 3 broods) for Ceriodaphnia showed no significant reductions at the three ambient stations in the second experiment when compared with the controls.

4.1.2 Contaminants Data

Inorganic contaminants data from the eight stations are presented in Table 13. Aluminum concentrations were reported above detection limits at all stations. Arsenic, tin and selenium were not reported above detection limits at any station. Concentrations of cadmium above detection limits were reported at Indian Head

The five Potomac U.S. EPA chronic water quality criteria for either freshwater or saltwater (U.S. Dahlgren. Indian Head, Freestone Point and Possum Point were freshwater stations; the other 5 stations were in saltwater. All underlined values exceed River stations are Indian Head, Freestone Point, Possum Point, Morgantown and Inorganic contaminants reports by date for the eight stations. EPA, 1987a). Table 13.

Station	Date	A1	As	cd	Cr	Contaminant Cu Hg	inant Hg	(ug/L) Ni	Pb	Sn	Se	Zn
Elizabeth R.	8/13/90	15.3	<6.0	<2.0	<3.0	3.2	<.2	<5.0	<2.0	<50	<30	14
	8/16/90	8.7	0.9>	<2.0	<3.0	3.7	0.22	<5.0	<2.0	<50	<30	15
Patapsco R.	8/13/90	20.3	<3.0	<2.0	<3.0	<2.0	<.2	10.8	<2.0	<50	<30	< 5
	8/16/90	5.4	<3.0	<2.0	<3.0	<2.0	<.2	6.5	<2.0	<50	<30	\ 5
Wye R.	8/13/90	10.9	<3.0	<2.0	<3.0	5.4	<.2	18	2.14	<50	<30	12.6
Indian Hd.	9/30/90	46.5	<3.0	1.32	<3.0	5.9	<.2	7.5	5.6	<5.0	<3.0	22.2
Freestone Pt.	9/27/90	49.8	<3.0	1.48	<3.0	6.7	<.2	12.0	5.6	<5.0	<3.0	26.5
Possum Pt.	9/27/90	48.6	<3.0	<1.0	<3.0	3.9	<.2	19.9	<2.0	<5.0	<3.0	19.6
Morgantown	9/24/90	10.3	<3.0	<1.0	<3.0	<2.0	• 29	8.4	2.8	<5.0	<3.0	<5.0
	9/27/90	16.8	<3.0	<1.0	<3.0	<2.0	<. 2 . 2	5.5	3.9	<5.0	<3.0	<5.0
Dahlgren	9/24/90	6.9	<3.0	<1.0	<3.0	<2.0	<.2	<5.0	5.2	<5.0	<3.0	<5.0
	9/27/90	15.2	<3.0	<1.0	<3.0	<2.0	<.2	5.5	3.6	<5.0	<3.0	7.4

(1.32 ug/L) and Freestone Point (1.48 ug/L). Copper concentrations above detection limits were reported from two Elizabeth River samples (3.2 and 3.7 ug/L), Wye River (5.4 ug/L), Indian Head (5.9 ug/L), Freestone Point (6.7 ug/L) and Possum Point (3.9 ug/L). Mercury was reported above detection limits from the Elizabeth River (0.22 ug/L) and Morgantown (0.29 ug/L) stations. Concentrations of nickel above detection limits were reported at all stations except the Elizabeth River and Dahlgren (9/24/90). Lead was reported above detection limits at the following stations: Wye River (2.14 ug/L), Indian Head (2.6 ug/L), Freestone Point (2.6 ug/L), Morgantown (2.8 and 3.9 ug/L) and Dahlgren (5.2 and 3.6 ug/L). Concentrations of zinc were reported above detection limits for all stations except the Patapsco River, Morgantown and one Dahlgren Sample.

Metal concentrations that exceeded EPA chronic water quality criteria for either freshwater (Indian Head, Freestone Point and Possum Point) or saltwater (all other stations) were: 1.32 and 1.48 ug/L cadmium at Indian Head and Freestone Point, respectively; 3.2 and 3.7 ug/L copper at the Elizabeth River station; 5.4 ug/L copper at the Wye River station; 0.22 and 0.29 ug/L mercury at the Elizabeth River and Morgantown stations, respectively, and 10.8, 18 and 8.4 ug/L nickel at the Patapsco River, Wye River and Morgantown stations, respectively (U.S. EPA, 1987a).

None of the organic contaminants listed in Table 2 were confirmed present above detection limits in any samples collected from the 8 stations. A minimum of one sample was analyzed from all

8 stations during testing. Two samples were analyzed from the Elizabeth River, Patapsco River, Potomac River - Morgantown and Potomac River - Dahlgren stations.

4.1.3 Water Quality Data

Water quality parameters reported from grab samples collected for all tests are presented in Table 14. The temperature and salinity (estuarine areas) were adjusted to 25 C and 15 ppt before exposing test species (except the ambient Elizabeth River salinity of 19-20 ppt). Most of these ambient water quality conditions appeared adequate for survival of test species with the possible exception of dissolved oxygen of 3.0 mg/L in the Elizabeth River.

Temperature, salinity, dissolved oxygen, pH and conductivity were measured at 1-h intervals at all Potomac River stations using datasonde units (Appendix B). All parameters appeared adequate for the various test species. Water quality parameters reported in test containers during tests are reported in Appendix C. Most of these parameters appeared adequate for test species except for occasional low D.O. values in the Elizabeth River test conditions.

4.1.4 Reference Toxicant Data

Twenty-four and 48-h LC50 values for the test species exposed to cadmium chloride during reference toxicant tests are presented in Table 15. Values obtained in these tests were compared with mean 48-h LC50 values for cadmium chloride collected over 2 years in our laboratory (except <u>E</u>. <u>affinis</u>). Forty-eight h LC50s for

Table 14. Water quality parameters reported in the field during collection of water samples. The following eight stations were sampled: Wye River (WR), Elizabeth River (ER), Patapsco River (PR), Morgantown (MT), Dahlgren (DG), Indian Head (IH), Possum Point (PP), Freestone Point (FP). Water quality parameters measured from the field samples collected from the Potomac River during the second Ceriodaphnia test are included.

Date 1990	Station	Temp (C)	Sal (ppt)	DO (mg/L)		Cond umhos/cm)	Hardness (mg/L CaCo ₃)
08/13	PR	24.0	7.5	8.8	8.58	12200	NA
	WR	27.0	9.0	7.4	8.13	15000	NA
	ER	28.7	20.1	3.3	7.22	32500	NA
08/16	PR	28.0	6.8	9.3	8.47	11600	NA
	WR	28.0	8.7	7.6	8.29	15200	NA
	ER	28.5	20.1	3.3	7.14	32300	NA
08/19	PR	27.8	6.3	7.2	8.52	11200	NA
	WR	27.0	9.5	5.0	8.28	16000	NA
	ER	28.2	19.4	3.0	7.24	31100	NA
09/24	DG	18.5	8.2	8.4	7.72	12000	NA
	MT	19.0	8.0	8.6	7.70	11800	NA
	PP	19.5	0	8.0	8.31	270	88
	IH	19.0	0	8.5	7.33	278	120
	FP	18.0	0	8.7	7.42	240	112
09/27	DG	18.0	8.0	8.2	-	12500	NA
	MT	19.0	7.5	7.4	7.46	11000	NA
	PP	19.5	0	8.0	-	268	100
	IH	20.5	0	7.8	_	283	108
	FP	19.0	0	9.3	-	255	92
09/30		21.0	8.0	8.8	-	13000	NA
	MT	21.0	8.5	7.4	_	13500	NA
	PP	20.0	0	7.0	-	285	112
	IH	20.2	0	6.6	-	203	124
	FP	20.0	0	8.0	-	305	100
10/26		14.5	0	10.5	7.75	200	92
	IH	14.0	0	10.0	7.60	150	92
	FP	14.0	0	10.8	7.77	200	84
10/28		13.0	0	8.8	7.80	173	92
	IH	12.5	0	10.0	7.69	140	76
	FΡ	13.0	0	9.0	7.69	160	84
10/31		12.0	0	9.8	7.55	160	80
	IH	12.5	0	9.4	7.57	140	84
	FP	11.0	0	10.0	7.69	173	90

NA = Not appropriate

Table 15. LC50's (mg/L) from reference toxicant tests conducted with cadmium chloride for the five species. Two reference toxicant tests were conducted with sheepshead minnow larvae because the source was different for the Potomac River experiments. The two-year mean 48-h value is based on numerous tests previously conducted in our laboratory.

Date 1990	<u>Species</u>	24-h	<u>48-h</u>	Two-year Mean 48 h	<u>96-h</u> *
07/18	Grass Shrimp	1.080	0.502	0.87	6. .:
07/30	Sheepshead Larvae	1.049	0.510	0.9	-
08/01	Eurytemora affinis	-	0.021		0.06
09/11	Fathead Larvae	0.123	0.117	0.09	-
09/17	Ceriodaphnia	0.449	0.134	0.076	-
10/20	Sheepshead Larvae	0.560	0.496	0.9	-

^{* 96-}h value from Roberts et. al, 1982

grass shrimp, fathead minnows, <u>Ceriodaphnia</u> and sheepshead minnows in the present study were similar to previous values. These data demonstrated that these species were healthy and test results from ambient toxicity tests were valid.

A 96-h LC50 value obtained for <u>E</u>. <u>affinis</u> nauplii from the literature was used for comparing our 48-h LC50 (Roberts et al., 1982). The 96-h LC value of 0.06 mg/L (95% confidence limits < .001 - .20 mg/L) reported by Roberts et al. (1982) was approximately one-third of our 48-h value (0.21 mg/L). The upper confidence limit of .20 mg/L reported by Roberts et al. (1982) was approximately equal to our 48-h values of .21 mg/L. Based on the 48-h difference in LC50 values, both of these concentrations are within the same range (i.e., the 96-h LC reported by Roberts et al. [1982] would likely be much higher after only 48 h). <u>E</u>. <u>affinis</u> cultures were therefore healthy, and test results from ambient tests were valid.

Two reference toxicant tests were conducted with sheepshead larvae because the sources of these fish were different. Both 48-h values were similar (0.510 mg/L and 0.496 mg/L).

4.2 <u>Sediment Tests</u>

The following results from sediment tests are presented below: toxicity data, contaminants data, and toxicity data from reference toxicant tests.

4.2.1 Toxicity Data

survival and growth data from toxicity tests at the three estuarine stations conducted on 8/28/90-9/17/90 in the Elizabeth River, Patapsco River, Wye River stations and controls are presented in Tables 16 and 17. Survival of all test organisms was statistically lower (0% survival) at the Elizabeth River station compared to the other stations and the controls. Survival of the grass shrimp was >99% at the other estuarine stations in this set.

None of the grass shrimp growth measures were statistically different from the controls.

The amphipod showed statistically lower survival in all three test sites compared to the controls (0% at the Elizabeth River, 54% survival at the Patapsco River, and 57% survival at the Wye River) after a 10 day exposure to test sediments. After 20 days exposure, the same trend was evident; significantly reduced survival in the Patapsco River (13% survival) and the Wye River (24% survival) stations as compared to the controls. Amphipod reburial data are presented in Table 18. The amphipod showed a significantly reduced ability to rebury in clean sediments after a 10 day exposure to the Patapsco River sediments. The amphipod also showed a reduced growth (in mean length) at the Wye River site as compared to the controls. The Wye River station was initially chosen as a "clean" reference site. The sediment sample from the Wye contained a large percentage of plant detrital material that may have interfered with the amphipods' ability to move within the sediments and feed, and therefore resulted in reduced survival and growth. In addition, the large detrital particles found in this sample may have

Table 16. Survival data from grass shrimp, amphipod and worm tests conducted at the Patapsco River, Elizabeth River and Wye River stations. Tests were conducted August 28, 1990 to September 17, 1990.

		% Surv	ival
<u>Species</u>	<u>Station</u>	<u>Day 10</u>	Day 20
Grass shrimp	Control	99	95
	Patapsco River	100	95
	Elizabeth River	0*	-
	Wye River	99	99
Amphipod	Control	98	83
	Patapsco River	54*	13*
	Elizabeth River	0*	- >
	Wye River	57*	24*
Worm	Control	100	100
	Patapsco River	58*	52*
	Elizabeth River	0*	-
	Wye River	46*	46*

^{*} Significantly different from the controls (p<0.05).

Table 17. Growth data (dry weight and length) for grass shrimp, amphipod and worm after 20-day exposure to sediments from the Patapsco River, Elizabeth River, and Wye River stations.

Site	Number of True Replicates*	Weight (mg)	S.E.	ength (mm)	S.E.
Grass shrimp:		(A)(C)			
Initial Meas	urements -	18.13	=	19.54	-
Control	5	32.56	2.20	23.74	0.65
Patapsco Riv	er 5	28.35	0.89	22.34	0.30
Wye River	5	27.37	1.12	22.73	0.45
Amphipod:					
Initial Meas	urements -	1.07	-	3.32	-
Control	5	1.10	0.08	4.44	0.17
Patapsco Riv	er 5	1.59	0.10	4.56	0.10
Wye River	5	1.13	0.14	4.05**	0.10
Worm:					
Initial Meas	surements -	0.54		6.25	-
Control	5	0.11	0.04	4.92	0.61
Patapsco Riv	ver 4	0.10	0.82	4.24	0.23
Wye River	3	0.14	0.02	5.08	0.70

^{*} Data for each replicate is the mean of the surviving animals from each.

^{**}Significantly different from the controls (p<0.05).

Table 18. Amphipod reburial data after 10 day exposure to sediments. Table shows percent of surviving animals able to rebury within one hour.

Station	<pre>% Reburial</pre>	<u>s.e.</u>
Patapsco River	<u>₹75*</u>	6.3
Wye River	95	3.6
Control	99	0.6
Dahlgren	100	0.0
Indian Head	100	0.0
Freestone Point	90	6.6
Possum Point	88	12.5
Morgantown	100	0.0
Nansemond Reference	98	15.3
Control	100	0.0

^{*} Significantly different from control (p<0.05).

interfered with retrieval of the animals at the end of the test period, resulting in seemingly high mortalities that may not have been totally due to toxicity.

The worms and amphipods showed similar results; statistically lower survival in the Patapsco River and the Wye River after 10 and 20 day exposures to test sediments, as compared to control survival. Survival in the Wye River sediments after 10 and 20 days of exposure was 46%. Survival of the worms in the Patapsco River sediments was 58% and 52% after 10 and 20 days, respectively. The worms showed no significant differences in growth, as compared to the controls.

Survival and growth data from estuarine and freshwater tests conducted at the five Potomac River stations and controls from 9/28/90-10/18/90 are presented in Tables 19 and 20. There was no significant difference in survival of the grass shrimp at any of the sites; all animals survived at all stations. There was a statistically significant reduction in growth (mean length) of the grass shrimp at the Possum Point site (19.90 mm) as compared to the controls (20.52 mm). None of the other shrimp measurements were statistically significant. The amphipod showed statistically reduced survival at all Potomac tests sites as compared to controls at both 10 and 20 day exposures, with only 6% survival at the Possum Point site after 10 days, and 0% survival after 20 days. No statistical differences were evident in the reburial data.

Worm survival was not statistically different at any of the Potomac River stations after 10 or 20-day exposures. Two

Table 19. Survival data from grass shrimp, amphipod and worm tests conducted at the five Potomac River stations. Tests were conducted September 28, 1990 to October 18,

<u>Species</u>		<u>Station</u>	% Surviv	val Day 20
Grass shrimp		Control	100	100
		Morgantown	100	100
		Dahlgren	100	100
		Possum Point	100	100
		Freestone Point	100	100
		Indian Head	100	100
		Nansemond Reference	e 100	100
Amphipod		Control	91	91
		Morgantown	70*	20*
		Dahlgren	22*	6*
		Possum Point	6*	0*
		Freestone Point	24*	4*
		Indian Head	41*	14*
		Nansemond Referenc	e 47*	12*
Worm		Control	74	57**
		Morgantown	61	55
		Dahlgren	61	62
		Possum Point	49	41
		Freestone Point	50	51
		Indian Head	67	53
		Nansemond Reference	e 51	40

^{*} Significantly different from the controls (p<0.05).

** Replicates with 0% survival due to predation were omitted from this calculation.

Table 20. Growth data (dry weight and length) for grass shrimp and amphipods after 20-day exposure to sediments from the five Potomac River stations.

	umber of True Replicates*	Weight (mg)		ength (mm)	S.E.
Grass shrimp:		Ba			
Initial Measu	res -	6.50	1	2.19	_
Dahlgren	5	24.86	1.43 2	20.71	0.16
Indian Head	5	23.09	1.47 2	20.56	0.37
Freestone Poin	nt 5	22.27	0.97 2	20.91	0.32
Possum Point	5	20.81	2.35	9.90**	0.38
Morgantown	5	21.13	1.48 2	21.21	0.25
Control	5	17.08	2.13 2	20.52	0.32
Nansemond Refe	erence 5	20.17	2.45 2	20.84	0.26
Amphipod:					
Initial Measu	res -	1.07	-	2.07	-
Dahlgren	2***	0.60	0.16	3.11	0.31
Indian Head	4	1.03	0.38	4.15	0.65
Freestone Poi	nt 3***	1.19	0.52	4.25	0.68
Possum Point	-		-	Ē	a-
Morgantown	4	0.90	0.10	4.08	0.14
Control	5	1.17	0.02	3.89	0.11
Nansemond Ref	erence 3	0.98	0.30	4.40	0.58

^{*} Data for each replicate is the mean of the surviving animals from each.

Note: Worm growth data were not collected due to sample preservation problems.

^{**} Significantly different from the controls (p<0.05).

^{***} Low survival (4-6%) may have reduced statistical sensitivity in detecting growth effects.

Streblospio benedicti at day 20. Several large predators (nereid worms) were also found in both test dishes. Control percent mortality was calculated omitting the two replicates with 0% survival. For future worm tests, all control sediments will be frozen \geq 48 hours prior to testing. No worm growth data were obtained for the Potomac stations because of a problem with the preservation technique used. All worms began to disintegrate before they could be measured.

4.2.2 <u>Contaminants Data</u>

Sediment samples from the eight stations were analyzed for Total Organic Carbon (TOC) and Acid Volatile Sulfides (AVS). results are shown in Table 21. The AVS approach to sediment contaminants evaluation is still developmental and only recently published (DiToro, 1990). To appropriately interpret the AVS data, selectively extractable metals (SEM) must be analyzed; a procedure that will be proposed for year two of this project. analysis of these samples will be used to help build a data base for future comparisons. TOC analysis was also included to allow for future comparisons. At present there is no readily accessible data base for comparison of TOC normalized data. Inorganic contaminants data from the eight stations are presented in Table 22. All test sites, the Nansemond Reference, and the worm control sediments had concentrations of the eleven metals above detection limits. At the Wye River station arsenic, cadmium and lead were

Table 21. Chemical data for sediment samples from the eight stations and the controls. The five Potomac River stations are Indian Head, Freestone Point, Possum Point, Morgantown, and Dahlgren. All data is on a dry weight basis.

<u>Station</u>	Total Organic Carbon (%)	AVS $(\mu \text{ mol/g})$
Elizabeth River (Atlantic Wood		5.90
Patapsco River	<0.37	4.60
Wye River	7.38	10.91
Amphipod Contro	<0.37	<1.00
Worm Control, S	et #1 1.54	5.02
Indian Head	2.77	5.32
Freestone Point	2.30	8.89
Possum Point	2.44	1.82
Morgantown	<0.37	<1.00
Dahlgren	0.39	3.15
Nansemond Refer	ence 1.42	4.03
Amphipod Contro Set #2	<0.37	<1.00
Worm Control, S	et #2 4.91	10.79

controls. The five Potomac River stations are Indian Head, Freestone Point, Possum concentrations exceeding "Effects Range-Low", and double underlined value represent concentrations exceeding "Effects Range-Medium" levels as defined in Long and Inorganic contaminants for sediment samples from the eight stations and the Point, Morgantown and Dahlgren. (Note: Single underlined values represent Morgan, 1990) Table 22.

Contaminant (ug/g)

Zn	275.7	603.1	6.8	29.9	207.7	211.9	171.1	223.6	31.0	40.8	2.0	246.7	149.4	0.1	120	270
Sn	15.0	83.0	3.0	7.0	8.0	18.0	14.0	19.0	5.0	5.0	<2.0	9.0	15.0	2.0	1	ı
Se	0.94	0.55	0.50	0.06	0.24	0.32	0.22	0.30	90.0	0.09	<0.02	0.26	0.39	0.02	1	ī
<mark>입</mark>	118.8	55.8	<3.8	14.5	118.2	47.3	40.2	47.8	6.6	10.5	<3.8	95.4	34.2	3.8	35.0	110.0
Ni	14.0	7.0	1.0	8.0	12.0	38.0	34.0	40.0	4.0	0.9	21.0	12.0	19.0	1.0	30	20
HG	3.226 14.0	0.022	0.075	<0.006	0.147 12.0	0.169	0.149 34.0	0.176 40.0	0.018 4.0	0.019 6.0	<0.00621.0	0.09912.0	0.09419.0	0.006	0.15	96.0
리	54.2	20.6	0.5	7.1	87.2	43.3	41.9	61.8	8.9	4.5	9.0	44.2	22.2	0.3	70.0	390.0
긺	30.0	157.0	2.0	11.0	29.0	45.0	47.0	48.0	22.0	11.0	12.0	29.0	39.0	1.0	80.0	145.0
AS	1.1	3.2	<0.3	0.4	1.0	1.0	0.7	0.7	9.0	0.3	<0.3	1.1	0.5	0.3	33.0	85.0
핑	23.28	4.89	<0.02	4.31	2.60	5.10	3.77	7.30	2.09	1.46	0.33	3,33	11.26	0.02	5.0	0.6
A1	12914.0	575.0	1100.0	7134.0	1005.0	19696.0	19409.0	23780.0	4593.0	3656.0	376.0	10954.0	19763.0	25.0	1	ı
Station	Elizabeth River	Patapsco River	Wye River	Amphipod Ctl*	Worm Ct1*	' Indian Head	Freestone Point	Possum Point	Morgantown	Dahlgren	Amphipod Ctl**	Worm Ct1**	Nansemond Ref.	Detection Limit	ER - L	ER - M

k = Set #1; ** = Set #2

all reported at concentrations below the detection limit. The amphipod control sediment used with the Potomac River test sediments had concentrations of cadmium, mercury, lead, selenium, and tin below detection limits.

Sediment-sorbed contaminants have been extensively studied by Long and Morgan (1990). They have established a table of concentrations at which biological effects would be expected if these contaminants were present in the sediment. The lower ten percentile of their data was established as the "Effects Range-Low" (ER-L) and median concentrations were identified as the" Effects Range-Median" (ER-M). Comparisons can be made between sites with respect to the potential for adverse biological effects by comparing the level of toxicants observed through chemical analysis with the ER-L or ER-M values. Those contaminants with levels exceeding the ER-L are in the "possible" effects range for toxic effects. The contaminant levels above the ER-M fall in the range of "probable" toxic effects.

The Elizabeth River station had concentrations of mercury, lead and zinc that all exceeded the ER-M as defined by Long and Morgan (1990). The Patapsco River site had concentrations of chromium (157.0 ug/g) and zinc (603.1 ug/g) that exceeded the ER-M (which is 145 ppm for chromium, and 270 ppm for zinc) and a concentration of lead (55.8 ug/g) that exceeds the ER-L of 35 ppm. Indian Head, Freestone Point, and Possum Point all had concentrations of mercury, nickel, lead and zinc that exceeded the ER-L for those contaminants (Table 22). The worm control used with the first set

(Elizabeth River, Patapsco River and Wye River) of tests had concentrations of copper, mercury, and zinc all above the ER-L and concentrations of lead above the ER-M. The worm control used with the Potomac stations also had concentrations of lead and zinc above the ER-L. The Nansemond Reference site had concentrations of zinc (149.4 ug/g) above the ER-L.

The sediment samples analyzed for semi-volatile organic compound and pesticides are presented in Appendix D and E. Only the Elizabeth River sediments had compounds listed in the Long and Morgan (1990) reference that exceeded the possible effects ranges. Nine of the sixteen semi-volatiles detected in the Elizabeth River sediments were above the ER-M values. In addition, two of the seven pesticides detected were above the ER-M values.

4.2.3 Pore Water Data

Sediment pore water was analyzed for sulfide, ammonia and nitrite for all eight stations and the controls and is presented in Table 23. Sediment samples used in toxicity tests were sieved with laboratory control water prior to conducting toxicity tests. The pore water analysis, however, was conducted on sediment samples collected from the field with no manipulation. The data from these analyses are included to provide relative numbers for comparison of test sites, rather than to suggest cause and effect relationships. It is assumed that the test pore water would reach a rough "steady state" with the sediments during the course of the experiments, so the pore water chemistry provides a relative indication of

Table 23. Chemical data for pore water samples from the eight stations and the controls. The five Potomac River stations are Indian Head, Freestone Point, Possum Point, Morgantown, and Dahlgren. All data is on a dry weight basis.

Station	Sulfide (mg/l)	Ammonia (mg/l)	Nitrite (mg/l)
Elizabeth Rive (Atlantic Woo		3.835	.0012
Patapsco Rive	r .013	2.099	.0049
Wye River Refe	erence .007	1.342	.0037
Amphipod Contr Set #1	rol, <.006	0.826	.0040
Worm Control, Set #1	.032	1.289	.0151
Indian Head	.035	6.477	.0070
Freestone Poin	nt .010	6.663	.0110
Possum Point	.012	15.790	.0205
Morgantown	.012	0.905	.0042
Dahlgren	.017	1.712	.0118
Nansemond Refe	erence .010	2.442	.0065
Amphipod Contr Set #2	rol, .042	2.489	.0028
Worm Control, Set #2	.023	3.176	.0034

potential stresses. Concentrations of ammonia greater than 6.0 mg/L were reported at Indian Head (6.5 mg/L), Freestone Point (6.7 mg/L) and Possum Point (15.8 mg/L). Levels of ammonia between 1.0 mg/L and 4.0 mg/L were reported at the Elizabeth River, Patapsco River, Dahlgren, Wye River, Nansemond Reference, the worm controls and amphipod control (set 2) sediments. The Morgantown sediments and the amphipod controls (set 1) had concentrations of NH3 < 1.0 mg/L. Sulfide concentrations at all test sites except Indian Head were below 0.02 mg/L. The Indian Head sediment pore water had a sulfide concentration of 0.035 mg/L. Control sediment sulfide concentrations ranged from <0.006 in the amphipod control (set 1) to 0.042 mg/L in the amphipod control, set 2.

4.2.4 Reference Toxicant Data

The relative sensitivities of each set of test organisms was evaluated by a reference toxicant test. The results of 96-hour static acute tests for grass shrimp exposed to sodium dodecyl sulfate (SDS) are presented in Table 24. Results of reference toxicant tests conducted with amphipods and worms exposed to cadmium chloride are also presented in Table 24. Although the grass shrimp tested were from two separate populations (one set reared at the AMRL and one set purchased), both sets of test animals showed similar sensitivities to the reference toxicant. Therefore, it is reasonable to consider toxicity data from the two sets of organisms valid and comparable. In addition, both tests gave results similar to those published in the literature. Tatem

Table 24. Reference toxicant data results from the ambient toxicity project.

Organism	Test Set #	Chemical	LC50 and CIs* (mg/l)
		8	
P. pugio	1	SDS**	56.7 (45.5 to 71.1)
	2	SDS	67.7 (54.6 to 83.9)
L. dytiscus	1	CdCl ₂	6.93 (5.25 to 9.26)
	2	CdCl ₂	5.58 (4.23 to 7.20)
S. benedicti	1	CdCl ₂	6.51 (5.12 to 8.21)
	2	CdCl ₂	5.69 (3.28 to 8.88)

^{*} CI = Confidence intervals **SDS = Sodium dodecyl sulfate

et al (1976) found median lethal toxicity values of 72.0, 90.0, 52.5, 98, and 55.0 mg/L SDS for grass shrimp held in the laboratory for less than a month. Values obtained in the amphipod test were comparable with 96-hr LC50 values for cadmium chloride collected over the last nine months in our laboratory. Amphipod reference toxicant tests conducted at the AMRL over a nine month period ranged from 6.13 mg/L CdCl₂ (95% CI 5.3 and 7.25) to 4.68 mg/L CdCl₂ (CI 3.70 and 6.09). The worm reference toxicant data represent the first set of tests done with this organism and are the beginning of a data base for future comparisons.

4.3 <u>Suborganismal Tests</u>

4.3.1 Cytochrome P-450 Induction

4.3.1.1. Fundulus Water and Sediment Exposures

Hepatic microsomal ethoxyresorufin-0-deethylase (EROD) activity in Fundulus exposed to test water and sediment is presented in Tables 25 and 26. Induction, or elevation, of EROD activity relative to controls or reference stations, suggests significant exposure to planar aromatic hydrocarbon inducing substances such as PAH and planar PCBs. Depression of activity is much less understood. High trace metal exposure can inhibit P-450 dependent activity via stimulation of heme degradation (cytochrome P-450 is a heme-protein). Alternatively, some organic substances can interfere with cytochrome P-450 metabolism through binding and inactivation of the catalyst (for example, insecticide synergists such as piperonyl butoxide) or through substrate competition at the

Table 25. Ethoxyresorufin-O-deethylase activity in <u>Fundulus</u> heteroclitus exposed to sediment and water from tests conducted 8/14/90 - 8/21/90.

Water Exposures	3	
Station	<u>(n)</u>	EROD Activity (pmol/min/mg protein)
Control	4	676.1 (327.3, 1396.4)
Wye River	5	379.3 (310.4, 463.4)
Patapsco R.	4	184.9 (105.9, 322.8) ^A
Elizabeth R.	2*	748.2 (140.0, 3999.4)
Sediment Expos	ures	
Control	5	323.8 ± 135.8 ⁸
Wye River	5	695.3 ± 442.3
Patapsco R.	6	404.4 ± 73.9
Elizabeth R.		Nearly Complete Mortality-No Samples.

^{*} Forty percent mortality. Mortality in other tests was less than 20%, except as noted for the Elizabeth River sediment exposure.

A Significantly different from the Control (p=0.05). To establish homogenous variance, an ANOVA was done on log transformed data and hence activities are reported as the backtransformed means with 95% confidence limits. Differences were tested with the Scheffe F Test (alpha=0.05).

⁸ Mean \pm SD.

Table 26. Ethoxyresorufin-O-deethylase activity in <u>Fundulus</u>
<u>heteroclitus</u> exposed to sediment and water from tests conducted 9/24/90 - 10/2/90 (Potomac River).

OD Activity (pmol/min/mg protein)
452.5 ± 164.2
571.2 ± 137.8
411.8 ± 148.9
401.3 ± 181.3
555.5 ± 147.1
471.9 ± 264.1
727.5 ± 518.4
617.6 ± 382.5
695.2 ± 278.1

Mortalities in all exposures were sporadic and less than 20%. All data as mean \pm s.d.

active site. This study is not designed to understand depression if it occurs, though the observation will be noted where appropriate. The appropriate comparisons for these data are within the exposure groups.

For water exposures, there were no significant differences detected between the controls and the stations tested (Dahlgren and Morgantown) in the estuarine portion of the Potomac River. There were, however, significant differences in EROD activity detected between animals from tests conducted with Wye, Patapsco and Elizabeth River waters. The most striking observation is the nearly 4-fold depression in activity in animals exposed to Patapsco River water. We did not detect any significant induction in activity between controls and test stations. It is important to note, however, that there was 40% mortality for Fundulus exposed to Elizabeth River water.

Mean values for EROD activity in sediment exposed <u>Fundulus</u> ranged approximately 2-fold and no significant differences were detected among stations from the two experiments. The variation encountered between the groups obscures any subtle differences between groups.

4.3.1.2 Larval Sheepshead and Fathead Minnows from Water Exposures

Samples of larval sheepshead and fathead minnows were collected for evaluation of monocygenase induction. These samples are currently being held at -80C until further processing.

4.3.1.3 Feral Fish

EROD activities were measured in samples of <u>Fundulus</u>, white perch and spot from various sampling locations (Table 27). Activities varied between species and the appropriate comparisons were made within a species among sites. Activities were highest in the spot samples, followed by <u>Fundulus</u> and white perch, respectively.

The <u>Fundulus</u> collected from the Wye River reference site and the Elizabeth River both contained high EROD activities. The activities were 2-5 fold higher than the mean activities detected in <u>Fundulus</u> from the water and sediment exposures (Tables 25 and 26).

White perch collected from four stations in the Potomac River had relatively low EROD activities and no significant differences were detected among stations. While we obtained samples from the Morgantown station, these samples have not been processed. We did note, however, that the liver samples from the largest white perch collected at this station were very dark brown in color. We will be processing these samples for histological evaluation and will attempt to resolve the cause of this phenomenon. From a consultation with Dr. Mike Lipsky of UMAB, we are hypothesizing that this "anomaly" may be due to high metal content. We recently discussed this observation with Dr. Tracie Bunton of Johns Hopkins University. She has been studying a copper storage disease in livers of white perch for the last several years. Her description of the color of livers of affected white perch ("looks like an old

Table 27. EROD activity in fish collected from various sampling locations

Species	(n)	EROD Activity (pmo	1/min/mg pr	cotein)
Fundulus heteroclitus				
Wye River	9	1584.8 ±	762.7	
Elizabeth River	15	2120.4 ±	1614.5	
White Perch				
Dahlgren	5	234.2 ±	133.0	
Possum Point	9	328.3 ±		
Indian Head	8	280.8 ±		
Freestone Pt.	9	233.9 ±		
Spot				
Wye River	3	2751.9 ±	634.3	
Patapsco River.	8	6315.2 ±		
Elizabeth River	6	5167.8 ±		
Freestone Pt.	3	2303.1 ±		

All data as mean ± s.d.

penny") agrees closely with our observations.

Spot were collected from four locations. EROD activity was high in all samples relative to the other species and the highest values were found in fish from the Elizabeth and Patapsco Rivers.

4.3.1.4 Ongoing Activities

We currently have many samples yet to process for the suborganismal component of year 1 studies. We have been processing samples in a logical fashion for six months and consider the results presented here as only preliminary. We will be conducting immunologic evaluations of the cytochrome P450IA concentration as the complement to the catalytic measurements. These studies will enable us to better resolve the extent of induction or repression of catalytic activities detected in the samples (see for example Gooch et al., 1989). We will also be evaluating the cytochrome P450 response in the larval fish samples collected from water exposures. In addition, we will also be carrying out laboratory studies with Fundulus using a model PAH inducer in order to calibrate the induction response in our local population of Fundulus.

In addition to completing cytochrome P-450 studies, we will be analyzing invertebrate samples for stress protein induction using immunoassay techniques with commercially available antibodies. As previously stated, it was apparent from the initiation of these studies, that the protocols found in the literature (i.e., the use of S-35 amino acids) would be logistically impossible for a study

of this size and design. We are, therefore, refocusing our efforts to determine the utility of this technique for year 2 studies and for ambient protocols in general.

SECTION 5

DISCUSSION

5.1 Elizabeth River Station

The Elizabeth River station located near the Atlantic Wood Industries (a wood treatment creosote facility) and in the vicinity of major military installations and industrial activities was known to be a toxic area based on previous data (Alden et al., 1988). Our water column toxicity tests confirmed the presence of toxic conditions. Significant reductions in survival were reported for E. affinis and grass shrimp after 8-d exposures to Elizabeth River water. Significant reductions in reproduction were not reported for surviving E. affinis and significant reductions in growth were not reported for surviving grass shrimp larvae. Survival and growth of sheepshead minnow larvae was not significantly reduced after 8-d exposures to water from this location.

This study was not designed to identify an absolute cause and effect relationship between biological effects and contaminant and/or water quality measurements (if present), due to the limited number and scope of these evaluations. Comparisons of field contaminant concentrations with specific water quality criteria (fresh or marine, acute or chronic) will be used throughout this discussion section to provide a basis for discussing the relationship between contaminant concentrations and biological

effects. These water quality criteria are based on toxicity data from numerous species (rather than a single species). It should be noted, however, that EPA water quality criterion were developed as values which are protective of most organisms and not necessarily concentrations at which stress or toxicity may occur. However, specific water quality criteria do provide a general "benchmark" for discussing possible effects if protective concentrations are exceeded.

At this particular station possibly stressful contaminant and water quality conditions were reported in the water column concurrently with significant reductions in survival of two test species. Copper concentrations of 3.2 and 3.7 ug/L in the Elizabeth River exceeded the U.S. EPA marine acute water quality criteria (2.9 ug/L) for this metal in saltwater (U.S. EPA, 1987a). In previous toxicity studies, concentrations of copper as low as 1 ug/L have been reported to inhibit the growth of phytoplankton Thalassiosira pseudonana (Davey et al., 1973). Copper concentrations at this station were higher than ambient concentrations of 0.08 - 2.3 ug/L reported in the main body of Chesapeake Bay waters (Kingston et al., 1983).

Mercury concentrations of 0.22 ug/L at this station also exceeded the U. S. EPA marine chronic water quality criteria (0.025 ug/L) for saltwater. It should be noted, however, that this mercury chronic criteria is based on human health data (e.g., FDA action levels and bioaccumulation data) and not aquatic toxicity data. Concentrations of total mercury lethal to sensitive aquatic

species range from 0.1 to 2.0 ug/L (Eisler, 1987). This extremely not be expected in detectable toxic heavy metal would concentrations in the ambient environment of Chesapeake Bay. Mercury is a particularly hazardous metal because non-toxic forms can be readily transformed into toxic forms (i.e., methylmercury) This metal can also be easily through biological processes. bioconcentrated in organisms and biomagnified through the food chain.

The above discussion of potentially toxic effects of copper and mercury at the Elizabeth River station should be interpreted with certain caveats. It is difficult to relate laboratory toxicity data with a field concentration of a specific metal without knowing the precise concentration of the toxic component of the metal that is bioavailable to the organism. Numerous physical (adsorption, flocculation sedimentation and remobilization), chemical (ionic redox reactions inorganic complexation, pH, strength, equilibria and organic-metallic compounds) and biological (uptake and incorporation, transformation and environmental modification) factors influence the bioavailability of these toxic metals. Although the above metals data were generated from .40 um filtered (dissolved fraction), we do not know the exact concentration of the toxic component of the bioavailable fraction.

Dissolved oxygen (~ 3.0 mg/L) was the only potentially stressful water quality condition reported from ambient water collected at this station. In laboratory studies, sensitive species such as larval stages of the American lobster (Homarus

americanus) were adversely affected by dissolved oxygen concentrations of ~3.0 mg/L (Don Miller, personal communication, U.S. Environmental Protection Agency). Jordan et al. (1990) have reported that dissolved oxygen concentrations of less than or equal to 3.0 mg/L can severely affect reproduction of Chesapeake Bay fish eggs and larvae. It is doubtful that the dissolved oxygen concentrations reported in the Elizabeth River were entirely responsible for the mortality of test species, but these conditions may have acted synergistically with other adverse contaminants.

The Elizabeth River station was located in a region with sediments previously demonstrated to be toxic (Alden and Young, 1982; Hargis et al., 1984; Alden and Butt, 1988; Alden et al., 1988) and highly contaminated with metals (Johnson and Villa, 1976; Alden et al., 1981; Rule, 1986; Alden et al., 1991) and organic pollutants (Alden and Hall, 1984; Bieri et al., 1986; Alden and Butt, 1988; Greaves, 1990; Alden et al., 1991). The in situ biological communities are clearly impacted by the contaminants in the sediments (Weeks and Warinner, 1986; Weeks et al., 1986; Ewing et al., 1989; Dauer et al., 1989; Alden et al., 1991). This station was selected to act as a "positive reference site", for which sediment toxicity tests should demonstrate significant effects.

All of the sediment test species (shrimp, worms and amphipods) exhibited complete mortality within the first 10 days of exposure to the sediments. The Elizabeth River sediment sample was 86.16% silt/clay by analysis. The amphipod Lepidactylus dytiscus will

exhibit approximately a 15% increase in mortality in clean sediments with greater than 85% silt/clay (Deaver and Adolphson, However, it was observed that all amphipods died in all replicates within the first few days of the test. Therefore, it is likely that the mortalities observed during the tests of sediments from this site are due to contaminants rather than particle size. The extremely high levels of contaminants in the sediments of the Elizabeth River station also indicated that this area is highly impacted. Mercury (3.23 ug/kg), lead (118.8 ug/g) and zinc (275.7 ug/kg) concentrations exceeded the ER-M levels employed by NOAA (Long and Morgan, 1990) to delineate sediments which are potentially toxic (Table 22). Each of these metals is potentially toxic to estuarine organisms (Waldichulk, 1974; Neff et al., 1978; Gilfillan et al., 1985; Long and Morgan, 1990) and could have been responsible for the observed toxicity. Unionized ammonia in pore water was also potentially toxic, with concentrations (0.36 mg/L) exceeding USEPA criteria for one-hour exposure (0.233 mg/L). However, potential organic toxicants, particularly polynuclear aromatic hydrocarbons (PNAHs), dominate the sediment contaminants at this site.

The sediments contained concentrations of PNAHs (Appendix D-7) that greatly exceeded almost all of ER-M values (9 out of 12) established for these contaminants by NOAA (Long and Morgan, 1990). Three additional semi-volatile organic contaminants (Appendix D-7) and over 20 tentatively identified semi-volatile contaminants (Appendix D-8) were detected in concentrations of thousands to tens

of thousands of parts per billion (ug/kg). In addition, seven pesticides were also detected, two of which exceeded the ER-M values (Appendix E-4). Previous studies have suggested that toxicity in the sediments from this region can be correlated with certain groups of PNAHs (Alden and Butt, 1987, 1988). In fact, reference toxicity tests suggest that fluoranthene in sediment alone is toxic to amphipods at concentrations below those observed for the Elizabeth River site (DeWitt et al., 1989; Deaver and Adolphson, 1991). Since fluoranthene is only one of numerous similar contaminants found at high concentrations at this station, the observed toxicity was not surprising. In fact, the magnitude of inorganic and organic contaminants found at highly elevated levels at this station would make sediment toxicity expected, but would confound speculations concerning specific "cause and effect" relationships.

The limited results from the suborganismal tests with <u>Fundulus</u> exposed to ambient water and sediment also suggested the presence of toxic conditions at this station. Forty percent mortality was reported in water exposures (at least double that of any other exposure) and <u>Fundulus</u> could not be maintained in the presence of Elizabeth River sediments. EROD activities in samples from surviving <u>Fundulus</u> from water exposure were not significantly elevated relative to the controls. However, native <u>Fundulus</u> collected from the Elizabeth River contained relatively high levels of EROD activity compared to other field studies with <u>Fundulus</u> (Elskus and Stegeman, 1989). Also, grossly visible nodules

(putatively tumors) were also noted in approximately 30% of the animals sampled from this site as has been seen by Vogelbein et al. (1990). This is not surprising in light of the high levels of PAH at this site.

The lack of apparent induction of EROD activity in <u>Fundulus</u> from water exposures indicates that either biologically available concentrations of PAH in the water were too low to cause induction (non-detectable in water samples at concentrations reported in Table 2) or that some other factor, chemical or biochemical, interfered with any induction response (note: we encountered 40% mortality in this group).

We also analyzed hepatic EROD activity in spot collected from this location. The values were approximately 2-fold higher than spot collected by Van Veld et al. (1990) from this location. It is also important to note that spot collected along a PAH contamination gradient in the Elizabeth River, did not show uniformly high values. Rather, hepatic EROD activity mirrored the gradient quite well. Thus, high EROD values in spot from this location were expected.

5.2 Patapsco River Station

Potentially toxic conditions were suspected at the Patapsco River station based on effluent toxicity data from a nearby industrial facility. However, actual toxicity data had not been previously collected at this specific study site. Results from our water column tests showed that survival of one of our three test

species (grass shrimp) was significantly reduced after 8 days of exposure to habitat water. Survival of <u>E. affinis</u> and sheepshead minnows was not significantly reduced in concurrent experiments. In addition, growth (sheepshead minnows) and reproduction (<u>E. affinis</u>) were not significantly reduced after exposure to Patapsco River water.

Contaminant and water quality data provided minimal information on potentially toxic conditions at the study site. Nickel concentrations of 10.8 ug/L at this station exceeded the U.S. EPA marine chronic water quality criteria for this metal in saltwater (U.S. EPA, 1987a). Therefore, concentrations of this metal may be stressful. All water quality conditions that were measured appeared adequate for the test species in water column tests.

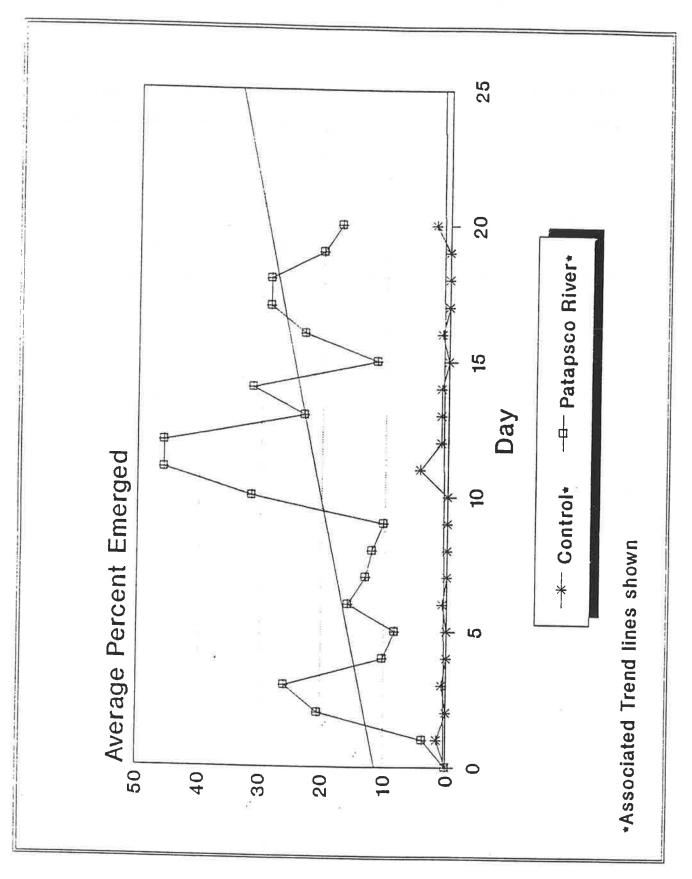
Sediment toxicity tests demonstrated toxic conditions at this site. Survival of two of the sediment test species, the amphipod and the worm, was significantly reduced after 10 and 20-d exposures to sediments collected from this site. Survival of the grass shrimp was not significantly reduced when compared to the controls. No significant growth effects were seen with any of the test organisms exposed to Patapsco River sediments. Although the amphipods showed no significant difference in growth at this station, they did show a significant difference in ability to rebury in clean sediments at the end of the 20-d test. Ninety-nine percent of the control animals were able to rebury, while only 75% of the surviving amphipods from the Patapsco River sediments were able to rebury when placed into clean sediment, indicating a

chronic level of toxicity (Table 18). In addition, observations during the test indicated that a high percentage of the organisms were emerged from the sediments or swimming in the overlying water throughout the test period. Figure 2 illustrates the average number of amphipods emerged per day throughout the 20-d test period as compared to the controls. This high percentage of emergence is abnormal for the amphipod Lepidactylus dytiscus, and indicates an avoidance response. In addition, it is possible that fewer mortalities may have occurred because the animals were swimming in the water column and not actually in contact with the sediment.

Very high levels of chromium, zinc, and lead were observed in the Patapsco River sediments. Chromium (157.0 ug/g) and zinc (603.1 ug/g) were found in levels exceeding the ER-M, and lead (55.8 ug/g) concentrations exceeded the ER-L level. The high levels of these metals is consistent with the various toxic responses exhibited at this site.

Suborganismal test results were similar to those reported for water column and sediment tests. Fundulus exposed to water samples from this location showed a significant depression in hepatic EROD activity as compared to controls. There are several possible explanations for this, though the results of catalytic protein measurements will be required for additional interpretation. Two logical hypotheses based on literature information are plausible. Previous studies have shown that high concentrations of PCBs can inhibit EROD activity in some teleosts (Gooch et al., 1989). It is possible, though unlikely, that this type of phenomenon could be

Figure 2. Amphipod emergence in Patapsco River and control sediments.



occurring here. An alternate possibility, is that high concentrations of trace metals could cause inhibition of activity, as has been seen with cadmium exposed flounder (George, 1989). Unfortunately, the contaminants data obtained do not support either of these hypotheses. Currently, we have no obvious explanation for these results.

In contrast, we measured high hepatic EROD activities in spot samples from this location. In fact, the values were similar to those found in spot collected from the Elizabeth River. These data are highly suggestive of biologically significant aromatic hydrocarbon exposure, putatively from somewhere near this location. This result is consistent with reported data on PNAH concentrations in sediments from in and around the Baltimore Harbor area (Helz and Huggett, 1987). Further analysis of the EROD catalyst content from this station, and others, will be required for a more detailed analysis of this observation.

5.3 Wye River Station

The Wye River station was selected for this study because it is located in a relatively pristine saltwater area where toxic conditions were not suspected. Survival, growth and reproduction data from our three test species suggested that habitat water from this station was non-toxic. It was interesting to note, however, that copper (5.4 ug/L) and nickel (18 ug/L) concentrations that exceeded the U.S. EPA water marine chronic quality criteria were reported at this station (U.S. EPA, 1987a). Toxic forms of these

metals (i.e., free cupric ion for copper) were likely not present, duration of exposure was too short to create toxic conditions, or reported concentrations were too low to cause effects with test species. All water quality conditions measured in the Wye River samples were adequate for survival, reproduction and growth of the test species.

In contrast to the water tests, sediment tests indicated potentially adverse conditions at the Wye River station. Significantly lower survival was reported in the amphipod and worm test. The amphipod had only 57% survival at the Wye station after 10 d of exposure and 24% survival after 20 d. Amphipod growth (mean length) was significantly lower than the controls. sediment collected at the Wye River station was 80% silt-clay and had a large percentage of organic plant matter (7.38% TOC). The combination of particle size and high organic matter may account, at least in part, for the reduced survival of the amphipod. DeWitt (1988) found that in fine, uncontaminated sediments (\geq 80% siltclay) mean survival of the amphipod Rhepoxynius abronius was approximately 15% lower than in native sediment. Deaver and Adolphson (1991) found similar results in tests done at the AMRL with the amphipod Lepidactylus dytiscus. In addition, the sediment sample from the Wye River site contained a high percentage of large detrital particles that may have interfered with the animals' ability to move and feed within the sediments and, therefore, may have resulted in reduced survival and growth.

The results of the worm bioassays were similar to those of the

amphipod tests. There was statistically reduced survival of the worms in the Wye River sediments after 10 and 20-day exposures. The worm showed no significant growth effects. Streblospio benedicti can occur naturally in areas of high organic matter and muddy sediments (Levin, 1981), therefore the sediment physical characteristics alone would not be expected to be the cause of the worm mortalities. The grass shrimp test did not suggest toxic conditions in Wye River sediment. Survival of grass shrimp was > 99% for both 10- and 20-d tests and no significant differences in growth were reported when compared with controls. Since two of the three species tested (the amphipod and the worm) showed significant reduction in survival at the Wye River station, it is possible that there was some unknown contaminant present that contributed to the toxic response. Very low concentrations of inorganic contaminants were reported in the sediment at this site. Unfortunately, organic contaminants were not analyzed due to funding limitations.

Suborganismal tests conducted at the Wye River site provided contrasting results for <u>Fundulus</u> tested in the laboratory versus feral fish. Hepatic EROD activity was not elevated relative to controls in <u>Fundulus</u> exposed to water and sediment from the Wye River. This potentially indicates no significant exposure to aromatic hydrocarbons through these routes or that other chemical or biological factors precluded induction (see discussion in Section 5.1). However, elevated EROD activities (relative to activities in sediment and water exposed animals) was reported in

Fundulus collected by seine from this Interestingly, the activities in the feral fish were near those detected in the fish collected from the Elizabeth River, a highly PNAH contaminated site. Since no information exists on possible genetic differences in P450 expression among Fundulus populations, a plausible hypothesis is that <u>Fundulus</u> from the Wye River are exposed to planar aromatic hydrocarbons, presumably through a dietary source. As previously discussed, organic contaminants (i.e., sediment concentrations of PNAHs or PCBs) were not measured, thus this suggestion cannot be corroborated. Spot collected from this site had mean hepatic EROD activities that were substantially lower than those collected from the Elizabeth River station, though unequal samples sizes and heterogeneous variance precludes a more thorough statistical evaluation of the feral spot data.

5.4 Potomac River - Indian Head Station

This station was selected because it is a representative freshwater area of the Potomac River near a major military facility with numerous outfalls. Although toxic conditions have not been previously documented at this station, results from our first Ceriodaphnia test demonstrated reduced survival. Contaminant and water quality data provide little insight on causes for reduced survival of Ceriodaphnia during experiment 1. Cadmium (1.32 ug/L) was the only metal measured at this station that was possibly toxic based on the U.S. EPA freshwater chronic water quality criterion of 1.1 ug/L (U.S. EPA, 1987a). All water quality conditions appeared

adequate for the test species. Results from one other freshwater test and three salinity adjusted tests showed no significant statistical reductions in survival, growth or reproduction for test species after exposure to ambient water from this station.

Sediment toxicity tests conducted at this station suggested toxic conditions. Results of the amphipod sediment tests indicated a significant reduction in survival at 10 and 20 days. This site was 88.82% silt/clay by analysis, which could account for approximately 15% mortality (Deaver and Adolphson, 1991). However, only 41% of the amphipods survived the initial 10-day test, and by day 20, only 14% of the test organisms remained alive. There were no significant effects seen in survival or growth with either the grass shrimp or the worms, or with growth of the amphipods. Inorganic contaminant analysis of the sediment indicated levels of mercury, nickel, lead and zinc that all exceeded the ER-L (see Table 22). Pore water analysis also showed high levels of ammonia (6.5 mg/L). By calculation, the level of unionized ammonia present at test conditions was 0.163 mg/L, which is near the EPA acute criteria level of 0.233 mg/L for a one hour average. The Indian Head sediments were not analyzed for organic contaminants.

Suborganismal tests (hepatic EROD activity) conducted to date at this station did not provide any data to suggest toxic conditions. Feral fish (white perch) collected at Indian Head had lower mean hepatic EROD values than spot from the Elizabeth and Patapsco Rivers.

5.5. Potomac River - Freestone Point

The Freestone Point station was located near an area where potentially toxic concentrations of trace metals (cadmium, chromium and copper) have been previously reported (Hall et al., 1987a). Survival, growth and reproduction data from our water column tests did not implicate the presence of toxic conditions at this site. Contaminant and water quality conditions from this station were generally non-toxic with the possible exception of cadmium. Cadmium concentrations of 1.48 ug/L at Freestone Point exceeded the U.S. EPA freshwater chronic water quality criteria of 1.1 ug/L for this trace metal in freshwater (U.S. EPA, 1987a). It is likely that the toxic form of cadmium was not present, duration of exposure was too short to create toxic conditions, or the reported concentration was too low to affect the test species.

One sediment toxicity test suggested toxic conditions at this station. Amphipods showed a significant decrease in survival (24% at day 10, 4% at day 20), but no growth effects. Again, the amphipod may have been somewhat affected by particle size (82.9% silt/clay) but only 15% of the deaths would be expected to be particle size preference (Deaver and Adolphson, 1991). The grass shrimp and worms showed no significant effects. Inorganic contaminants analysis indicated levels of mercury, nickel, lead and zinc that exceeded the ER-L (see Table 22). Pore water analysis showed ammonia levels of 6.663 mg/L and organic analysis identified two semi-volatile compounds (Appendix D-3), twenty tentatively identified semi-volatiles (Appendix D-4) and two pesticides (Appendix E-2) present.

Suborganismal tests conducted to date suggest no significant toxic conditions based on laboratory exposures of <u>Fundulus</u> to sediment and evaluation of EROD activity in feral fish (white perch and spot).

5.6 Potomac River - Possum Point

This station was located in an area where potentially toxic concentrations of trace metals have been reported in previous striped bass contaminant studies (Hall et al., 1990; Hall, unpublished data). Survival, growth and reproduction data from our water column tests did not show toxic conditions at this station. Water quality and contaminants data from this station also suggested that the ambient water from this station was non-toxic.

Results from sediment toxicity tests indicated a significant decrease in survival of the amphipod at day 10 and day 20. Only 6% of the test animals were alive after 10 days of exposure to Possum Point sediments, and none were alive after 20 days of exposure. The Possum Point sediments were 98% silt/clay by analysis, which could account for up to 15% of the amphipod mortalities at this site. The grass shrimp and the worm showed no significant differences in survival, but there was a significant difference in grass shrimp mean length, as compared to the controls. Inorganic analysis indicated high concentration of mercury, nickel, lead and zinc. Concentrations of these metals exceeded the ER-L. In addition, pore water analysis indicated very high levels of ammonia. By calculation, the level of unionized ammonia present

(0.316 mg/L) under test conditions would have exceeded the EPA acute criteria level of 0.233 mg/L for a one hour average. Organic analysis identified five semi-volatiles (Appendix D-1), 20 tentatively identified semi-volatiles (Appendix D-2) and the pesticide DDE present in the sediment sample (Appendix E-1). Any of these toxicants, or a combination of effects from all of them, may be the source of the toxicity in this sediment sample. The response of the amphipod illustrates the varying sensitivities of different organisms, and serves to underscore the need for multispecies toxicity testing.

Suborganismal tests (hepatic EROD activity) conducted to date at this station did not suggest the presence of toxic conditions. Feral fish (white perch) collected at Possum Point did not demonstrate any significant induction of hepatic EROD activities.

5.7 Potomac River - Morgantown

The Morgantown station was located in a typical mesohaline area of the Potomac River. Toxic conditions have not been previously reported at this station with the exception of tributyltin concentrations of 20 - 24 ng/L in 1985 and 1986 (Hall et al., 1987b). Results from one of our water column tests suggested toxic conditions were present. Survival of sheepshead minnow larvae was significantly reduced (40%) at this station after 8 days of exposure when compared with control survival (88%). Survival and reproduction of <u>E. affinis</u> and survival and growth of grass shrimp was not significantly reduced at this station. The sheepshead

minnow is generally considered to be a moderately resistent species; therefore, reduced survival reported at this station is noteworthy. Survival of test species in our most toxic area (the Elizabeth River) showed no reduction in survival of sheepshead minnow larvae but reduced survival was reported for our other two test species (E. affinis and grass shrimp). These data support the use of water column tests with several species when attempting to evaluate toxic conditions in ambient areas.

Contaminants data from this station showed that concentrations of mercury (0.29 ug/L) and nickel (8.4 ug/L) exceeded the U.S. EPA marine chronic water quality criteria for these metals in saltwater (U.S. EPA, 1987a). Mercury is an extremely toxic heavy metal as concentrations of 0.1 ug/L have been reported toxic to sensitive aquatic biota (Eisler, 1987). Nickel is much less toxic than mercury but the 8.4 ug/L still exceeds environmentally safe concentrations as determined by the U.S. Environmental Protection Agency. All water quality conditions measured in the water column appeared non-toxic.

Results from the amphipod sediment toxicity test showed a statistically significant reduction in survival (70% survival at day 10 and 20% survival at day 20). The Morgantown sediment was approximately 92% sand, very similar to the native amphipod sediment, so the differences in mortality cannot be attributed to sand-silt/clay ratios. No toxic levels of inorganic contaminants or unionized ammonia were found in this sediment. Since organic analysis was not conducted on this sample due to funding

limitations, it is possible that the amphipod mortalities are due to an unknown organic contaminant. The shrimp and the worm showed no significant difference in survival or growth.

Suborganismal tests conducted to date at this station did not suggest the presence of toxic conditions based on exposures of Fundulus to water and sediment. Feral white perch collected from the Morgantown station did provide interesting anecdotal information as their livers were dark brown (see Section 4.3.1.3).

5.8 Potomac River - Dahlgren Station

This station was located in a typical mesohaline area of the Potomac River near a major military facility. Toxic conditions have not been previously evaluated in this area. Results from our sheepshead minnow larvae tests showed reduced survival of larvae after 8 days of exposure. Survival, growth and reproduction data from the E. affinis and grass shrimp tests showed no significant effects at this station. Contaminant and water quality data showed no obvious conditions that were potentially toxic based on U.S. EPA water quality criteria data or toxicity data with sheepshead minnow larvae. It is interesting to note, however, that detectable concentrations of lead (3.6 and 5.2 ug/L) were reported at this station.

Results from sediment toxicity tests indicated a significant reduction in amphipod survival (22% survival), but no significant differences in shrimp and worm survival or shrimp growth.

Contaminant analysis showed no evidence of toxic inorganic

contaminants. The sediments from this site were > 85% sand by analysis, similar to native amphipod sediment. Therefore, the mortalities seen at this site would not be due to particle size preference of the amphipods. Pore water analysis did show relatively high levels of unionized ammonia (0.159 mg/L), just below the EPA acute criteria of 0.233 mg/L for a one hour average of unionized ammonia. The amphipod appears to be more sensitive to ammonia stress than the other sediment species tested. Organic analysis identified two semi-volatiles (Appendix D-5), 11 tentatively identified semi-volatile compounds (Appendix D-6), and one pesticide (Appendix E-3) present in the sample.

Suborganismal tests conducted to date do not suggest any significant induction of hepatic EROD activity in fish from this site. The results for feral fish (white perch) were similar.

SECTION 6

CONCLUSIONS

The primary goal of this pilot study was to use a suite of water column, sediment and suborganismal tests to identify toxic areas in the Chesapeake Bay watershed. This battery of tests demonstrated the presence of toxic conditions in various environmental media in suspected contaminated areas such as the Elizabeth River and the Patapsco River (Table 28). importantly however, some of these tests suggested the presence of toxic conditions in a suspected toxic-free habitat in the Wye River and in critical habitat areas of the Potomac River (Table 28). Results from this study clearly demonstrate the need multispecies testing within each type of test when attempting to identify toxic ambient areas. Two water column tests (\underline{E} . affinis and sheepshead minnows) conducted at the Patapsco River station showed no significant effects. However, the third test species (larval grass shrimp) demonstrated reduced survival at this station. The same scenario existed for sediment tests conducted at all Potomac River sites. The worm tests showed no effects at all 5 stations and the grass shrimp demonstrated no effects at 4 of the stations. In contrast, reduced survival of amphipods was reported at all 5 stations.

The need for integrated water column, sediment and suborganismal testing was also demonstrated because one type of

Patapsco River; WR = Wye River; IH = Indian Head: FP = Freestone Pt; PP = Possum Effects were NE = No Effect; M = Mortality; RG = Reduced Growth; RR = Reduced Reproduction; RB = Reduced Reburial; IE = Induced EROD Activity; RE = Reduced EROD Activity and LA = Liver Abnormality. Station abbreviations were as follows: ER = Elizabeth River; PR noted if significantly different than controls for water column and sediment Toxicity data from water column, sediment and suborganismal tests. tests. The following abbreviations were used for various effects: Pt; MT = Morgantown and DA = Dahlgren. Table 28.

	ER	PR	WR	Station IH	FP	PP	MT	DA
Test Type Water Column	Later Control of the							ļ
E. affinis P. pugio C. variegatus Ceriodaphnia sp.	EE E I	NE*	N N N I I	NE* NE NE M, NE	N N N N N N N N N N N N N N N N N N N	N N N N N N N N N N N N N N N N N N N	X X X I I	* E E E I I
Sediment								ā
Amphipod Worm Grass Shrimp	N N N	M, RB M NE	M, RG NE	M NE NE	ж ы ы ы	NE RG	N N E E	N N S
Suborganismal**							;	
<u>Fundulus</u> - water <u>Fundulus</u> - sediment Feral Fish	M M IE	NE IE	NE NE IE	NE NE	NE NE	N N I	NE LA	N N N

o Fo *Low control reproduction limits the use of this endpoint; therefore, NE is based survival.

**Qualitative determinations.

test was not sufficient to maximize our ability to identify toxic conditions in the ambient environment of the Chesapeake Bay watershed. For example, the three water column tests conducted at the Wye River station did not suggest the presence of toxic conditions but the sediment and suborganismal tests did implicate potential contaminant problems. Sediment tests with amphipods also identified toxic conditions at two Potomac River stations (Freestone Point and Possum Point) where five water column tests did not. This integrated multi-test approach also demonstrated supportive data for the various stations. The water column and sediment tests demonstrated significant effects at the Elizabeth River and Patapsco River stations; suborganismal tests suggested significant effects at these stations. Both the water column and sediment tests also showed significant effects at the Indian Head, Morgantown and Dahlgren stations in the Potomac River.

Ranking of sensitivity among tests within each test type (water column, sediment and suborganismal) showed the following results after one year of testing. Results from the water column tests demonstrated no significant ranking of sensitivity among the three saltwater tests but rather supported the need for multispecies tests because different species display varying sensitivity to different types of contaminants. Both grass shrimp and sheepshead minnow tests demonstrated effects at two of the eight stations, while <u>E</u>. affinis showed effects at only one station. A comparison of the two freshwater water column tests at three stations showed that the <u>Ceriodaphnia</u> test was more sensitive than the fathead

minnow test. Results from the sediment tests clearly showed that the amphipod test was most sensitive as effects were reported at all stations. The polychaete worm test was the second most sensitive test and the grass shrimp test was least sensitive. A ranking of sensitivity among the various suborganismal tests is not appropriate until further research has been completed.

Water quality and contaminant evaluations conducted in water and sediment during this pilot study provided supportive information on possible causes of biological effects but these evaluations were not intended to provide conclusive data on specific cause and effect relationships if reported. The number of potentially toxic inorganic contaminants in water and inorganic and organic contaminants in sediment was higher at the station where the greatest number of toxic effects were reported by biological indicators (Elizabeth River). The second most toxic area based on biological data, the Patapsco River, also contained potentially toxic concentrations of inorganic contaminants in both water and sediment. The Wye River station provided an interesting comparison between biological and contaminants data. Both sediment and suborganismal tests suggested toxic conditions while water column tests did not. Toxic inorganics were not detected in Wye River sediment and organic evaluations were not conducted in this media. In contrast, two potentially toxic metals were reported in the water column at the Wye River station.

The Potomac River was perhaps the most important habitat evaluated in this pilot study. It provided various examples where

toxic effects demonstrated in our tests were reported concurrently with potentially toxic contaminants. There are also examples of biological effects without supportive contaminants data. One potentially toxic metal was reported in the water column at the Indian Head and Freestone Point station and four potentially toxic metals were measured in the sediment at each of these stations. Results from one water column test and one sediment test demonstrated toxic conditions at Indian Head. However, only one sediment test showed toxic conditions at Freestone Potentially toxic metals were not found in the water column at Possum Point but four potentially toxic metals were found in the The amphipod test (sediment exposure) demonstrated reduced survival while no significant effects were reported during Potentially water column or suborganismal tests. contaminants were not reported in sediment from the Morgantown or Two metals exceeded U.S. EPA water quality Dahlgren station. criteria at Morgantown and no potentially toxic contaminants were reported in the water column at Dahlgren. Significant effects were reported from water column and sediment tests at both Morgantown and Dahlgren.

SECTION 7

REFERENCES

- Alden, R.W., III and A.J. Butt. 1988. Toxicity testing of dredged materials using multiple-species microcosms. <u>In</u> Chemical and Biological Characterization of Sludges, Sediments, Dredge Spoils, and Drilling Muds (J.J. Lichtenberg, F.A. Winter, C.I. Weber and L. Fradkin, eds.) pp 418-437. Amer. Soc. for Testing and Materials, Philadelphia, PA.
- Alden, R.W., III and A.J. Butt. 1987. Statistical classification of the toxicity and polynuclear aromatic hydrocarbon contamination of sediments from a highly industrialized seaport. Environm. Toxicol. Chem. 6:673-684.
- Alden, R.W., III, A.J. Butt and R.J. Young, Jr. 1988. Toxicity testing of sublethal effects of dredged materials. Arch. Environm. Contam. and Toxicol. 17:381-389.
- Alden, R.W., III., A.S. Gordon, E.F. Stillwell, R.K. Everton and M.F. Helmstetter. 1988. An evaluation of the distribution of toxicants/mutagens in the Elizabeth River, Virginia, in relation to land use activities. Final report to the Virginia Water Control Board.
- Alden, R.W., III, D.M. Dauer and J.H. Rule. 1981. An assessment of the ecological impact of open ocean disposal of materials dredged from a highly industrialized estuary. Final Report to the National Oceanic and Atmospheric Administration. 472 pp.
- Alden, R.W., III., R.M. Ewing and M.F. Helmstetter. 1990.

- Distribution and management implications of contaminants in Elizabeth River sediments: I. Metal and organic contaminants and total organic carbon. Draft final reported to the Virginia Water Control Board.
- Alden, R.W., III, R.M. Ewing, M.F. Helmstetter, J.G. Winfield, D.J. Burdige, J.H. Rule, G.T.F. Wong, D.J.P. Swift and A. Muller. 1988. Distribution and management implications of contaminants in Elizabeth River sediments: A compilation of recent research on sediment contamination in the Elizabeth River. Final Report to the Virginia State Water Control Board, Richmond, VA. 263 pp., 4 Appendices.
- Alden, R.W., III., A.S. Gordon, E.F. Stillwell, R.K. Everton and M.F. Helmstetter. 1988. An evaluation of the distribution of toxicants/mutagens in the Elizabeth River, Virginia, in relation to land use activities. Final report to the Virginia Water Control Board.
- Alden, R.W., III and G.J. Hall. 1984. Polynuclear aromatic hydrocarbons and toxicity of sediments from the Elizabeth River, Hampton Roads, Virginia. Supplemental contract report to the U.S. Army Corps of Engineers, Norfolk District. 52 pp.
- Alden, R.W., III and R.J. Young, Jr. 1982. Open ocean disposal of materials dredged from a highly industrialized estuary: An evaluation of potential lethal effects. Arch. Environm.

 Contam. and Toxicol. 11:567-576.
- APHA et al. (American Public Health Association, American Water Associates of Water Pollution Control Federation). 1989.

- Standard Methods for the Examination of Water and Wastewater, 17th Ed. American Public Health Association, Washington, DC.
- APHA et al. (American Public Health Association, American Water Works Association, Water Pollution Control Federation). 1985.

 Standard Methods for the Examination of Water and Wastewater.

 American Public Health Association, Washington, DC.
- ASTM, 1990. Guide for Conducting 10-day Static Sediment Toxicity

 Tests with Marine and Estuarine Amphipods. Standard E-1367.

 Janet Lamberson and Richard Swartz (eds.) ASTM, Philadelphia,
 PA.
- Bender, M.E. and R. J. Huggett. 1987. Contaminant effects on Chesapeake Bay Shellfish. <u>In</u> Contaminant Problems and Management of Living Chesapeake Bay Resources. (S.K. Majumdar, L.W. Hall, Jr. and H.M. Austin, eds) pp 373-393. Pennsylvania Academy of Sciences, Easton, PA.
- Bieri, R.H., C. Hein, R.J. Huggett, P. Shou, H. Slone, C. Smith and Chih-Wu Su. 1986. Polycyclic aromatic hydrocarbons in surface sediments from the Elizabeth River subestuary. Int. J. Environm. and Anal. Chem. 26:97-113.
- Bradley, B.P., R. Hakimzadeh and J.S. Vincent. 1988. Rapid response to stress in <u>Eurytemora affinis</u>. Hydrobiol. 167/168:197-200.
- CEC (Chesapeake Executive Council). 1988a. Chesapeake Bay basinwide toxics reduction strategy. Chesapeake Bay Agreement Commitment Report. Chesapeake Bay Liaison Office, Annapolis, MD.

- CEC (Chesapeake Executive Council). 1988b. Chesapeake Bay living resource monitoring plan. Chesapeake Bay Agreement Commitment Report. Chesapeake Bay Liaison Office, Annapolis, MD.
- Chapman, P.M. 1988. Marine Sediment Toxicity Test. <u>In</u> Chemical and Biological Characterization of Sludges, Sediments, Dredge Spoils and Drilling Muds. ASTM STP 976 (J.J. Lichtenberg, F.A. Winter, C.I. Weber and L. Fradkin, eds.) pp 391-402. American Society for Testing and Materials, Philadelphia, PA.
- Dauer, D.M., H.G. Marshall, R.S. Birdsong, R.M. Ewing and A.J. Rodi. 1989. Elizabeth River long-term monitoring program:

 Biological monitoring projects. Final Report for Phase I to Virginia State Water Control Board, Richmond, VA. 66 pp.
- Davey, E.W., M.J. Morgan and S.J. Erickson. 1973. A biological measurement of the copper complexation capacity of seawater. Limnol. and Oceanog. 18:993-997.
- Deaver, E. and P.C. Adolphson. 1991. Evaluation of the amphipod

 Lepidactylus dytiscus as a sediment toxicity test organism.

 (Manuscript in review).
- DeWitt, T.H., G.R. Dittsworth and R.C. Swartz. 1988. Effects of
 Natural Sediment Features on Surviving the Phoxocephalid
 Amphipod. Rheopoxynius abronius. Mar. Environ. Res. 25:99124.
- DeWitt, T.H., R.C. Swartz and J.O. Lamberson. 1989. Measuring the acute toxicity of estuarine sediments. Environ. Toxicol. Chem. 8:1035-1048.
- DiToro, D.M., J.D. Mahony, D.J. Hansen, K.J. Scott, M.B. Hicks,

- S.M. Mayr and M.S. Redmond. 1990. Toxicity of cadmium in sediment: the role of acid volatile sulfide. Environ. Tox. and Chem. 9:1489-1504.
- Elskus, A.A. and J.J. Stegeman. 1989. Induced cytochrome P-450 in Fundulus heteroclitus associated with environmental contamination by polychlorinated biphenyls and polynuclear aromatic hydrocarbons. Mar. Environ. Res. 27:31-50.
- Eisler, R. 1987. Mercury hazards to fish, wildlife and invertebrates: A synoptic review. Fish and Wildlife Service Biological Report 85 (1.10), Laurel, MD.
- Ewing, R.M., R.W. Alden, III., S.W. Sokolowski and E. Deaver.

 1989. Elizabeth River long-term monitoring program, Phase I:

 Water quality monitoring, contaminants in water and sediments
 and toxicity tests. Interim report to the Virginia Water

 Control Board.
- Ewing, R.M., R.W. Alden, III, S.W.Sokolowski and E. Deaver. 1989.

 Elizabeth River long-term monitoring program: Water Quality

 monitoring contaminants in water and sediments and toxicity

 tests. Final Report for Phase I to Virginia State Water

 Control Board, Richmond, VA. 56 pp.
- Fisher, D.J., D.T. Burton, L.W. Hall, Jr., R.L. Paulson and C.M. Hersh. 1988. Standard operating procedures for short-term chronic effluent toxicity tests with freshwater and saltwater organisms. Johns Hopkins University, Applied Physics Laboratory, Aquatic Ecology Section, Shady Side, MD.
- George, S.G. 1989. Cadmium effects on plaice liver xenobiotic and

- metal detoxication systems: dose-response. Aquat. Toxicol. 15:303-309.
- Gilfillan, E.S., D.S. Page, D. Vallas, L. Gonzalez, E. Pendergast, J.C. Foster and S.A. Hanson. 1985. Relationship between glucose-6-phosphate dehydrogenases and aspartate amino transferase activities, scope for growth, and body burdens of Ag, Cd, Cu, Cr, Pb and Zn in populations of Mytilus edulis from a polluted estuary. In Marine Pollution and Physiology: Recent Advances. (F.J. Vernberg, F.P. Thurberg, A. Calabrese and W.B. Vernberg, eds.) pp 11-30. University of South Carolina Press.
- Gooch, J.W., A.A. Elskus, P.J. Kloepper-Sams, M.E. Hahn and J.J. Stegeman. 1989. Effects of ortho- and non-ortho-substituted polychlorinated biphenyl congeners on the hepatic monooxygenase system in scup (Stenotomus chrysops). Toxicol. Appl. Pharmacol. 98:422-433.
- Greaves, J. 1990. Elizabeth River long-term monitoring program,

 Phase I: Analysis of organic pollutants in sediments and blue

 crab (<u>Callinectus sapidus</u>) tissues. VIMS Final report to the

 Virginia Water Control Board, Richmond, VA.
- Guillard, R.R.L. 1975. Culture of phytoplankton for feeding marine invertebrates. <u>In</u> Culture of Marine Invertebrate Animals (W.L. Smith and M.H. Chanley, eds.) pp 29-60. Pleum Publishing, New York, NY.
- Hall, L.W. Jr., S.J. Bushong, W.S. Hall and W.E. Johnson. 1988c.

 Acute and chronic effects of tributyltin on a Chesapeake Bay

- copepod. Environ. Toxicol. Chem. 7: 41:46.
- Hall, L.W. Jr., S.J. Bushong, M.C. Ziegenfuss and W.S. Hall.

 1988b. Concurrent mobile on-site and in-situ striped bass
 contaminant and water quality studies in the Choptank River
 and Upper Chesapeake Bay. Environ. Toxicol. and Chem. 7: 815830.
- Hall, L.W. Jr., W.S. Hall, S.J. Bushong and R.L. Herman. 1987a.

 In-situ striped bass, <u>Morone saxatilis</u>, contaminant and water quality studies in the Potomac River. Aquatic Toxicol. 10:-73
- Hall, L.W. Jr., M.J. Lenkevich, W.S. Hall, A.E. Pinkney and S.J. Bushong. 1987b. Evaluation of butyltin compounds in Maryland waters of Chesapeake Bay. Marine. Pollut. Bulletin 18: 78-83.
- Hall, L.W. Jr., A.E. Pinkney, L.O. Horseman and S.E. Finger. 1985.

 Mortality of striped bass larvae in relation to contaminants
 and water quality in a Chesapeake Bay tributary. Trans. Am.
 Fish Soc. 114: 861-868.
- Hall, L.W. Jr., M.C. Ziegenfuss, S.J. Bushong, J.A. Sullivan and M.A. Unger. 1990. Striped bass contaminant and water quality studies in the Potomac River and Upper Chesapeake Bay in 1989: Annual contaminant and water quality evaluations in east coast striped bass habitats. Report. U.S. Fish and Wildlife Service, Leetown, WV.
- Hall, L.W. Jr., M.C. Ziegenfuss, S.J. Bushong and M.A. Unger.

 1988a. Striped bass contaminant and water quality studies in
 the Potomac River and Upper Chesapeake Bay: Annual

- contaminant and water quality evaluations in east coast striped bass habitats. Report. The Johns Hopkins University Applied Physics Laboratory, Aquatic Ecology Section, Shady Side, MD.
- Hall, L.W. Jr., M.C. Ziegenfuss, S.J. Bushong, M.A. Unger and R.L. Herman. 1989. Studies of contaminant and water quality effects on striped bass prolarvae and yearlings in the Potomac River and Upper Chesapeake Bay in 1988. Trans. Am. Fish. Soc. 188: 619-629.
- Hargis, W.J., M.H. Roberts and D.E. Swerner. 1984. Effects of contaminated sediments and sediment-exposed effluent water on an estuarine fish: Acute toxicity. Mar. Environm. Res. 14:337-354.
- Heikkila, J.J., G.A. Schultz, K. Iatrou and L. Gedamu. 1982.

 Expression of a set of fish genes following heat or metal ion exposure. J. Biol. Chem. 257:12000-12005.
- Heinle, D.R. and D.A. Flemmer. 1975. Carbon requirements of a population of the estuarine copepod, <u>Eurytemora affinis</u>.

 Marine Biology. 31: 235-247.
- Helz, G.R. and R.J. Huggett. 1987. Contaminants in Chesapeake Bay: The regional perspective. <u>In</u> Contaminant Problems and Management of Chesapeake Bay Living Resources. (S.K. Majumdar, L.W. Hall, Jr. and H.M. Austin, eds.) pp 270-297. Pennsylvania Academy of Sciences, Easton, PA.
- Jimenez, B.D. and J.J. Stegeman. 1990. Detoxication enzymes as indicators of environmental stress on fish. Am. Fish. Soc.

- Symp. 8:67-69.
- Johnson, P.G. and O. Villa, Jr. 1976. Distribution of metals in the Elizabeth River. EPA Annapolis Field Office Tech. Rep. 61. 903/9-76-023.
- Jordan, S., K. Mountford, C. Stenger, R. Batiuk, M. Olson and D. Forsell. 1990. Chesapeake Bay dissolved oxygen restoration goals. Report. Maryland Department of Natural Resources, Annapolis, MD.
- Kingston, H.M., R.R. Greenberg, E.S. Beary, B.R. Moody, T.C. Rains and W.S. Liggett. 1983. The characterization of Chesapeake Bay: A systematic analysis of toxic trace elements. U.S. Department of Commerce, National Bureau of Standards, Washington, DC.
- Klauda, R.J. and M.E. Bender. 1987. Contaminant effects on Chesapeake finfishes. <u>In</u> Contaminant Problems and Management of Chesapeake Bay Living Resources. (S.K. Majumdar, L.W. Hall, Jr. and H.M. Austin, eds.) pp 373-393. Pennsylvania Academy of Sciences, Easton, PA.
- Klauda, R.J., R.E. Palmer and D.M. Baudler. 1988. Effects of acidic episodes in Lyons Creek, Maryland, on survival of the eggs and larvae of blueback herring and American shad: 1987 field and laboratory studies. The Johns Hopkins University Applied Physics Laboratory, Shady Side, MD.
- Kleinow, K.M., M.J. Melancon and J.J. Lech. 1987.

 Biotransformation and induction: Implications for toxicity,

 bioaccumulation and monitoring of environmental xenobiotics in

- fish. Environ. Health Persp. 71:105-119.
- Kloepper-Sams, P.J., S.S. Park, H.V. Gelboin and J.J. Stegeman.

 1987. Specificity and cross-reactivity of monoclonal and polyclonal antibodies against cytochrome P-450E of the marine fish scup. Arch. Biochem. Biophys. 253:268-278.
- Kloepper-Sams, P.J. and J.J. Stegeman. 1989. The temporal relationships between cytochrome C-450E protein content, catalytic activity and m-RNA levels in the teleost, <u>Fundulus heteroclitus</u> following treatment with B-naphoflorane. Arch. Biochem. Biophy. 268:525-535.
- Levin, L.A. 1981. Dispersion, feeding behavior and competition in two spinoid polychaetes. J. Mar. Res. 39(1):99-117.
- Long, E.R. and L.G. Morgan. 1990. The Potential for Biological Effects of Sediment-Sorbed Contaminants Tested in the National Status and Trends Program. National Technical Memorandum Nos. OMA 52. Seattle, WA.
- Masters, J.A., M.A. Lewis, D.H. Davidson and R.D. Bruce. 1991.

 Validation of a four-day <u>Ceriodaphnia</u> toxicity test and statistical consideration in data analysis. Environ. Toxicol.

 Chem. 10:47-55.
- Neff, J.W., R.S. Foster and J.F. Slowey. 1978. Availability of sediment-adsorbed heavy metals to benthos with particular emphasis on deposit feeding infauna. Tech. Rep. D-78-42.

 U.S. Army Corps of Engineers Waterways Exp. Sta., Vicksburg, MS.
- Roberts, M.H., J.E. Warinner, C.F. Tsai, D. Wright and L.E. Cronin.

- 1982. Comparison of estuarine species sensitivities to three toxicants. Arch. Environ. Contam. Toxicol. 11: 681-692.
- Roccheri, M.C., M.G. DiBernardo and G. Giudice. 1981. Synthesis of heat-shock proteins in developing sea urchins. Dev. Biol. 83:173-177.
- Rule, J.H. 1986. Assessment of trace element geochemistry of Hampton Road Harbor and lower Chesapeake Bay area sediments. Environ. Geol. Water Sci. 8(4):209-219.
- Sanders, B.M. 1988. The role of the stress protein response in physiological adaptation of marine molluscs. Mar. Env. Res. 24:207-210.
- Scientific Instruments Inc. 1981. Ammonia in water and seawater.

 SIC Method No. 116-D223-01, Scientific Instruments Corp.,

 Hawthorne, NY.
- Stegeman, J.J. 1989. Cytochrome P450 forms in fish: catalytic immunological and sequence similarities. Xenobiotica. 19(10):1093-1110.
- Stegeman, J.J., F.Y. Teng and E.A. Snowberger. 1987. Induced cytochrome P450 in winter flounder (Pseudopleuronectes americanus) from coastal Massachusetts evaluated by catalytic assay and monoclonal antibody probes. Can. J. Fish Aquat. Sci. 44:1270-1277.
- Stephan, C.E. 1978. LC50 TOXDAT Program. Unpublished Data.

 Environmental Research Laboratory, U.S. Environmental

 Protection Agency, Duluth, MN.
- Swartz, R.C., W.A. DeBen, J.L.P. Jones, J.O. Lamberson and F.A.

- Cole. 1985. Phoxocephalid amphipod bioassay for marine sediment toxicity. In Aquatic Toxicology and Hazard Assessment: Seventh Symposium (R.D. Cardwell, R. Purdy and R.C. Bahner, eds.) pp 287-307. STP 854. American Society for Testing and Materials, Philadelphia, PA.
- Tatem, H.E., J.W. Anderson and J.M. Neff. 1976. Seasonal and laboratory variations in the health of grass shrimp Palaemonetes pugio: dodecyl sodium sulfate bioassay. Bull. Environ. Contam. Toxicol. 16:368-375.
- Towbin, H., T. Staehelin and J. Gordon. 1979. Electrophoretic transfer of proteins from polyacrylamide gels to nitrocellulose sheets: procedure and some applications.

 Proc. Natl. Acad. Sci. USA. 76:4350-4354.
- U.S. EPA (United States Environmental Protection Agency). 1987a.
 Water quality criteria summary. U.S. Environmental Protection
 Agency Office of Water Regulations and Standards. Criteria and
 Standards Division, Washington, DC.
- U.S. EPA (United States Environmental Protection Agency). 1987b.
 Ambient aquatic life water quality advisories for tributyltin.
 Narragansett, RI.
- U.S. EPA (United States Environmental Protection Agency). 1986.

 Test Methods for Evaluating Solid Waste Laboratory Manual Physical/Chemical Methods. USEPA SW-846. Washington, DC.
- U.S. EPA (United States Environmental Protection Agency). 1985a.

 Technical support document for water quality-based toxics control. Office of Water Enforcements and Permits and Office

- of Water Regulations and Standards, Washington, DC.
- U.S. EPA (United States Environmental Protection Agency). 1985b.

 Short-term methods for estimating the chronic toxicity of effluents and receiving waters to freshwater organisms.

 Environmental Monitoring and Support Laboratory. EPA 600/4-85-014. U.S. EPA, Cincinnati, OH.
- U.S. EPA (United States Environmental Protection Agency). 1985.

 Methods for Measuring the Acute Toxicity of Effluents to

 Freshwater and Marine Organisms, 3rd ed. (W.H. Peltier and

 C.I. Webers, eds.) pp 216. EPA-600/4-85-013. USEPA,

 Cincinnati, OH.
- U.S. EPA (United States Environmental Protection Agency). 1984.

 Guidelines establishing test procedures for the analysis of pollutants under the Clean Water Act. Federal Register 49 (209): 43234-43442.
- U.S. EPA (United States Environmental Protection Agency). 1984.

 Final water quality standards. Fed. Reg. 48:51400-51443.
- U.S. EPA (United States Environmental Protection Agency). 1983.

 Final water quality standards. Fed. Reg. 48: 51400-51443.
- U.S. EPA (United States Environmental Protection Agency). 1979.
 Methods for chemical analysis of water and wastes. EPA 600/4-79-020. U.S. EPA, Cincinnati, OH.
- USEPA/ACOE. 1977. Ecological Evaluation of Proposed Discharge of Dredged Material into Ocean Waters. <u>In Implementation Manual</u> for Section 103 of Public Law 92-532 (Marine Protection, Research, and Sanctuaries Act of 1972). Technical Committee

- on Criteria for Dredged and Fill material. July 1977 (second printing April 1978) Environmental Effects Lab, U.S. Army Corps of Engineers Waterways Experiment Station, Vicksburg, MS.
- Van Veld, P.A., J.J. Stegeman, B.R. Woodin, J.S. Patton and R.F. Lee. 1988. Induction of monooxygenase activity in the intestine of spot (<u>Leiostomus xanthurus</u>), a marine teleost, by dietary polycyclic aromatic hydrocarbons. Drug Metab. Dispos. 16:659-665.
- Van Veld, P.A., D.J. Westbrook, B.R. Woodin, R.C. Hale, C.L. Smith, R.J. Huggett and J.J. Stegeman. 1990. Induced cytochrome P-450 in intestine and liver of spot (<u>Leiostomus xanthurus</u>) from a polycylic aromatic hydrocarbon contaminated environment.

 Aquat. Toxicol. 17:119-132.
- Vogelbein, W.K., J.W. Fournic, P.A. Van Veld and R.J. Huggett.

 1990. Hepatic neoplasms in the mummichog (<u>Fundulus</u>

 <u>heteroclitus</u>) from a creosote contaminated site. Cancer Res.

 50:5978-5986.
- Waldichuk, M. 1974. Some biological concerns in heavy metals pollution. <u>In</u> Pollution and Physiology of Marine Organisms (F.J. Vernberg, ed.) Academic Press, NY.
- Weber, C.I., et al. 1989. Short-term methods for estimating the chronic toxicity of effluents and receiving waters to freshwater organisms, second edition. Environmental Monitoring Systems Laboratory, U.S. Environmental Protection Agency, Cincinnati, OH, EPA/600/4-89/001.

- Weeks, B.A. and J.E. Warinner. 1986. Functional evaluation of macrophages in fish from a polluted estuary. Vet. Immunol. Immunopathol. 12:313-320.
- Weeks, B.A., J.E. Warinner, P.L. Mason and D.S. McGinnis. 1986.

 Influence of toxic chemicals on the chemotactic response of fish macrophages. J. Fish Biology 28(6):653-658.
- Wright, D.A., C.H. Hocutt and I. Hartwell. 1988. Bioassessment of pollutant stress in Chesapeake Bay organisms. Interim Progress Report. University of Maryland, Chesapeake Biological Laboratory, Solomons, MD.

Appendix A

Suborganismal sampling for various fish species

SAMPLING RECORD FOR AMBIENT TOX. STUDY- FISH

				VIAL	NO.	TISSU		BODY	GONAD	
SPECIES	TEST TYPE	SITE	DATE	NO.	POOLED	LIV	GUT	WEIGHT	WEIGHT	SEX
SPOT	- Feral	BEAR CR.	8/12/90	1	1	×	X	32.3	<.01	
SPOT	Feral	BEAR CR.	8/12/90	2	1	X	X	13.7	<.01	
SPOT	Feral	BEAR CR.	8/12/90	3	1	X	X	25.7	0.02	
SPOT	Feral	BEAR CR.	8/12/90	4	1	X	X	18.3	0.01	
SPOT	Feral	BEAR CR.	8/12/90	5	1	X	X	48.8	<.01	
SPOT	Feral	BEAR CR.	8/12/90	6	1	X	X	39	<.01	
SPOT	Feral	BEAR CR.	8/12/90	7	1	X	X	15.2	<.01	
SPOT	Feral	BEAR CR.	8/12/90	8	1	X	X	49.3	0.04	
SPOT	Feral	BEAR CR.	8/12/90	9	1	X	X	26.1	0.02	
SPOT	Feral	BEAR CR.	8/12/90	10	1	X	X	36.9	0.01	
SPOT	Feral	BEAR CR.	8/12/90	11	1	X	X	30.7	<.01	
SPOT	Feral	BEAR CR.	8/12/90	12	1	X	X	23.6	0.03	
SPOT	Feral	ELIZABETH	8/27/90	1	1	X	X	32.7	0.06	
SPOT	Ferai	ELIZABETH	8/27/90	2	1	X	X	15.2	<.01	
SPOT	Ferai	ELIZABETH	8/27/90	3	1	X	X	17.4	0.02	
SPOT	Ferai	ELIZABETH	8/27/90	4	1	X	X	25.6	0.05	
SPOT	Feral	ELIZABETH	8/27/90	5	1	X	X	22.2	<.01	
SPOT	Feral	ELIZABETH	8/27/90	6	1	X	X	33.8	<.01	
SPOT	Feral	ELIZABETH	8/27/90	7	1	X	X	15.3	0.02	
SPOT	Feral	ELIZABETH	8/27/90	8	2	X	X	26.5	0.02	
SPOT	Feral	ELIZABETH	8/27/90		1	X	X	18.1	<.01	
FUNDULU		ELIZABETH	8/16/90		1	X	X	14	0.34	F
FUNDULU		ELIZABETH	8/16/90		1	X	X	11.1	0.63	F
FUNDULU		ELIZABETH	8/16/90		1	X	X	6.9	0.16	F
FUNDULU		ELIZABETH	8/16/90		1	X	X	12.7	0.28	M
FUNDULU		ELIZABETH	8/16/90		1	X	X	8.4	0.26	M
FUNDULU		ELIZABETH	8/16/90		1	X	X	6.2	0.06	М
FUNDULU		ELIZABETH	8/16/90		1	X	X	7		F
FUNDULL		ELIZABETH	8/16/90		1	X	X	6.4	0.14	M
FUNDULL		ELIZABETH	8/16/90		1	X	X	11.8	0.21	M
FUNDULL		ELIZABETH	8/16/90		1	X	X	5.5	0.32	F
FUNDULL		ELIZABETH	8/16/90		1	X	X	6	0.04	M
	JS Feral	ELIZABETH	8/16/90		1	X	X	8.9	0.34	F
	JS Ferai	ELIZABETH	8/16/90		1	X	X	6.7	0.17	F
	JS Feral	ELIZABETH	8/16/90		1	X		7.1	0.22	F
	JS Feral	ELIZABETH	8/16/90		1	Х		9	<.01	M
	JS Feral	ELIZABETH	8/16/90		1	Х		6.5	0.05	M
	JS Feral	ELIZABETH	8/16/90		1	X		10.9	0.14	M
	JS Feral	ELIZABETH	8/16/90			X		8.6	0.11	M
	JS Ferai	ELIZABETH	8/16/90		1	X		6.2	0.07	M
	JS Feral	ELIZABETH	8/16/90		1	X		6.7	0.08	М
	US Feral	WYER	8/15/90		1	X		15.9	0.27	F
	US Feral	WER	8/15/90		i i	X				M
	US Feral	WYER	8/15/90		1	X			0.33	F
	US Feral	WER	8/15/9		- 1	X			0.05	
	US Feral	WER	8/15/9		1	X			0.06	
CONDOL	US FEIRI	AAICU	011019	, ,	'	^			2.50	

FU	NDULUS	Feral	WYER	8/15/90	6		1	X	X	11.9	0.07	М
	NDULUS		WYER	8/15/90	7		1	X	Χ	14.2		F
	NDULUS		WER	8/15/90	8		1	X	X	6.4	0.01	M
	INDULUS		WYER	8/15/90	9		1	X	Χ	24.1	0.91	F
	INDULUS		WYER	8/15/90	10		1	X	X	11.4		F
	INDULUS		WYER	8/15/90	11		1	X	X	16.1	0.08	M
	INDULUS		WER	8/15/90	12		1	X	X	10.9	0.01	M
	INDULUS		WYER	8/15/90	13		1	X	X	5.5	<.01	M
	INDULUS		WER	8/15/90	14		1	X	X	5.3	0.01	М
	INDULUS		WER	8/15/90	15		1	X	X	9.1	0.05	М
	INDULUS		WER	8/15/90	16		1	X	X	6.9	0.3	F
	INDULUS		WYER	8/15/90	17		i	×	X	5.8	0.03	F
	UNDULUS		WER	8/15/90	18		i	X	X	5.4		F
	JNDULUS		WER	8/15/90	19		1	x	,.	9.3	0.04	F
	INDULUS		WER	8/15/90	20		i .	x		5.3	0.02	F
			WER	8/15/90	1		1	x		22.4	0.11	F
	MAJELIS			8/15/90	2		1	x		23.2	0.15	F
	MAJELIS		WER				1	x		17.5	0.08	F
	MAJELIS		WER	8/15/90	3		-	x		14.9	0.07	F
	MAJELIS		WER	8/15/90	4		1		Х	32.8	0.07	•
	POT	Feral	WYER	8/15/90	1		1	X	x	52.6	0.05	
	POT	Feral	WYER	8/15/90	2		1	X	X	23.7	<.01	
	POT	Feral	WYER	8/15/90	3		1	X	Α.			
	POT	Feral	WYER	8/15/90	4		1	X		40	<.01	
	.BASS	Ferai	WYER	8/15/90	1		1	X		55.2	0.02	
	.BASS	Feral	WYER	8/15/90	2		1	X		67.8	0.13	
	.BASS	Ferai	WYER	8/15/90	3		1	Х		59.6	<.01	
W	/. PERCH	Ferai	DAHLGREN	9/26/90	1		1	X	X	28.2	0.31	
V	/. PERCH	Feral	DAHLGREN	9/26/90	2		1	X	X	26.2	0.03	
V	/. PERCH	Feral	DAHLGREN	9/26/90	3		1	X	X	30.3	0.05	
V	V. PERCH	Feral	DAHLGREN	9/26/90	4		1	X	X	23.9	0.16	
V	V. PERCH	Feral	DAHLGREN	9/26/90	5		1	X	Χ	16.6	0.11	
٧	V. PERCH	Feral	DAHLGREN	9/26/90	6		1	X	Х	15.4	0.1	
٧	V. PERCH	Feral	DAHLGREN	9/26/90	7		2	X		25.4	0.14	
٧	V. PERCH	Feral	DAHLGREN	9/26/90	8		2	X		27.9	0.07	
٧	V. PERCH	Ferai	DAHLGREN	9/26/90	9		2	X		29	0.1	
٧	V. PERCH	Feral	DAHLGREN	9/26/90	10		3	X		38.2	0.07	
٧	V. PERCH	Ferai	MORGANTN	9/26/90	1		1	X	X	44.4	0.29	
٧	V. PERCH	Feral	MORGANTN	9/26/90	2		1	X	X	51.1	0.22	
٧	V. PERCH	Feral	MORGANTN	9/26/90	3		1	X	X	41.2	0.24	
٧	V. PERCH	Feral	MORGANTN	9/26/90	4		1	X	X	38.6	0.33	
٧	V. PERCH	Feral	MORGANTN	9/26/90	5		1	X	X	48.2	0.08	
٧	V. PERCH	Feral	MORGANTN	9/26/90	6		1	X	X	52.1	0.46	
	V. PERCH		MORGANTN	9/26/90	7		1	X		37.5	0.22	
	V. PERCH		MORGANTN	9/26/90	120 8		3	Χ		55.2	0.05	
	V. PERCH		POSSUM PT	9/26/90	1		1	X	X	34.9	0.26	
	V. PERCH		POSSUM PT	9/26/90	2	97	1	X	X	37.5	0.32	
	V. PERCH		POSSUM PT	9/26/90	3		1	X	X	37	0.29	
	V. PERCH		POSSUM PT	9/26/90	4		1	X	Χ	28.2	0.12	
	V. PERCH		POSSUM PT		5		1	X	X	25.5	0.04	
	V. PERCH		POSSUM PT	9/26/90	6		1	X	X	34.9	0.26	
	W. PERCH		POSSUM PT	9/26/90	7		2	X		58.2	0.28	
	W. PERCH		POSSUM PT	9/26/90	8		3	X		95.3	0.35	
	W. PERCH		POSSUM PT	9/26/90	9		2	X		69.9	0.48	
,		r c rai	F COOUNT F I	3120130	J		~	^		50.0	3	

								0.40	
W. PERCH Fera		9/26/90	10	2	X		56.5	0.12	
W. PERCH Fera	I FREESTONE	9/26/90	1	1	X	X	24	0.15	
W. PERCH Fera	I FREESTONE	9/26/90	2	3	X	X	40.4	0.26	
W.PERCH Fera	FREESTONE	9/27/90	1 "	1	X	X	223	2.11	
W. PERCH Fera	I FREESTONE	9/27/90	2	1	X	X	6 6 .4	0.46	
W. PERCH Fera	I FREESTONE	9/27/90	3	1	X	Χ	34.6	0.2	
W. PERCH Fera	I FREESTONE	9/27/90	4	1	X	X	31.9	0.1	
W. PERCH Fera	i FREESTONE	9/27/90	5	1	X	X	34.7	0.12	
W. PERCH Fera		9/27/90	6	1	X	X	22.4	0.01	
W. PERCH Fera		9/27/90	7	2	X		46.4	0.2	
W.PERCH Fera		9/27/90	8	2	X		40.9	0.22	
W.PERCH Fera		9/27/90	1	1	X	X	41.9	0.11	
W.PERCH Fera		9/27/90	2	1	X	X	37.2	0.23	
W.PERCH Fera		9/27/90	3	1	X	X	43.2	0.34	
W.PERCH Fera		9/27/90	4	1	X	X	33.2	0.24	
W.PERCH Fera		9/27/90	5	1	X	X	33.3	0.23	
W.PERCH Fera		9/27/90	6	1	X	X	31.8	0.03	
W.PERCH Fera		9/27/90	7	1	X		48.4	0.48	
		9/27/90	8	1	X		40.2	0.12	
		9/27/90	9	2	X		62.2	0.28	
W.PERCH Fera			10	2	X		62.9	0.42	
W. PERCH Fera		9/27/90		3	x		56.5	0.03	
SPOT Fera		9/27/90	1		x		41.4	0.05	
SPOT Fera		9/27/90	1	2			46.6	0.02	
SPOT Fera		9/27/90	2	2	X		32	0.02	
SPOT Fera		9/27/90	3	2	X			0.04	M
FUNDULUS WAT		10/2/90	1	1	X		7.5		
FUNDULUS WAT		10/2/90	2	1	X		6.3	0.05	F
FUNDULUS WAT		10/2/90	3	1	Х		5	0.12	
FUNDULUS WAT		10/2/90	4	1	X		6.6	0.1	F
FUNDULUS WA		10/2/90	5	1	X		5.4	0.08	М
FUNDULUS WA		10/2/90	6	1	X		5.9	0.04	F
FUNDULUS WA	TER CONTROL	10/2/90	7	1	X		4.7	0.03	M
FUNDULUS WA	TER CONTROL	10/2/90	8	1	X		3.9	0.03	F
FUNDULUS WA	TER CONTROL	10/2/90	9	1	X		5.4	0.03	F
FUNDULUS WA	TER CONTROL	10/2/90	10	1	X		3.5	0.01	M
FUNDULUS WA	TER CONTROL	10/2/90	11	1	X		3.5	0.02	F
FUNDULUS WA	TER CONTROL	10/2/90	12	1	X		4.8	0.05	F
FUNDULUS WA	TER CONTROL	10/2/90	13	4			16.8		
FUNDULUS WA	TER MORGANTN	10/2/90	1	1	X		9.4	0.13	M
FUNDULUS WA		10/2/90	2	1	X		7.4	0.38	F
FUNDULUS WA		10/2/90		1	Х		7.2	0.04	F
FUNDULUS WA		10/2/90		1	Х		5.7	0.02	М
FUNDULUS WA		10/2/90		-1	Х		9.5	0.24	F
FUNDULUS WA				1	Х		6	0.02	M
FUNDULUS WA				1	X		7.2	0.05	F
FUNDULUS WA		10/2/90			X		5.5	0.05	F
FUNDULUS WA			•	1	X		7.1	0.05	M
FUNDULUS WA				1	X		4.9	0.02	М
FUNDULUS WA				1	X		4.6	0.03	М
FUNDULUS WA				<u> </u>	x		4.8	0.01	М
				4	^		17.5		
FUNDULUS WA				1	Х		11.4	0.11	М
FUNDULUS WA		10/2/90			X		11.6	0.04	М
FUNDULUS WA	NTER DAHLGREN	10/2/90	2	1	^		11.0	3.07	

FUNDULUS '	WATER	DAHLGREN	10/2/90	3	1	X	6.1	0.01	
FUNDULUS '			10/2/90	4	1	X	7.4	0.08	F
FUNDULUS '		_	10/2/90	5	1	X	5.3	0.05	F
FUNDULUS			10/2/90	6	1	X	5.4	0.01	M
FUNDULUS		DAHLGREN	10/2/90	7	1	Х	5.1	0.01	M
FUNDULUS		DAHLGREN	10/2/90	8	1	X	5	0.04	F
FUNDULUS		DAHLGREN	10/2/90	9	1	X	6.1	0.04	F
FUNDULUS		DAHLGREN	10/2/90	10	i	X	3.4	0.05	F
FUNDULUS		DAHLGREN	10/2/90	11	1	X	4.2	0.04	F
				12	1	X	5.9	0.06	F
FUNDULUS		DAHLGREN	10/2/90		3	x	11.7	0.00	•
FUNDULUS		DAHLGREN	10/2/90	13	1	x	4.5	0.08	F
FUNDULUS		CONTROL	8/21/90	1	-	x	12.5	0.19	F
FUNDULUS		CONTROL	8/21/90	2	1		2.9	0.13	F
FUNDULUS		CONTROL	8/21/90	3	1	X		<.01	M
FUNDULUS		CONTROL	8/21/90	4	1	X	2.1		F
FUNDULUS		CONTROL	8/21/90	5	1	X	7.4	0.06	
FUNDULUS		CONTROL	8/21/90	6	1	X	4.3	<.01	M
FUNDULUS	WATER	CONTROL	8/21/90	7	1	X	5.9	0.21	F
FUNDULUS	WATER	CONTROL	8/21/90	8	1	X	7	0.02	M
FUNDULUS	WATER	CONTROL	8/21/90	9	1	X	7	0.05	F
FUNDULUS	WATER	CONTROL	8/21/90	10	1	X	4.4	0.02	F
FUNDULUS	WATER	WYER	8/21/90	1	1	X	5.1	0.07	F
FUNDULUS	WATER	WYER	8/21/90	2	1	X	9.3	0.06	F
FUNDULUS	WATER	WYER	8/21/90	3	1	X	2.9	<.01	М
FUNDULUS	WATER	WYER	8/21/90	4	1	X	10.9	0.05	M
FUNDULUS		WER	8/21/90	5	1	X	10.5	0.08	F
FUNDULUS		WER	8/21/90	6	1	X	6.3	0.09	F
FUNDULUS		WYER	8/21/90	7	1	X	4.3	0.01	M
FUNDULUS		WYER	8/21/90	8	1	X	6.2	0.05	F
FUNDULUS		WER	8/21/90	9	2	X	4.6	0.01	М
FUNDULUS		BALTIMORE	8/21/90	1	1	X	4.6	0.04	F
FUNDULUS		BALTIMORE	8/21/90	2	i	X	9	0.09	F
		BALTIMORE	8/21/90	3	i	X	6.8	0.02	M
FUNDULUS					1	X	4.6	0.04	M
FUNDULUS		BALTIMORE	8/21/90	4	1	x	5.9	0.05	F
FUNDULUS		BALTIMORE	8/21/90	5	=	x	6.9	0.02	M
FUNDULUS		BALTIMORE	8/21/90	6	2		6.1	0.02	F
FUNDULUS		BALTIMORE	8/21/90	7	2	X	6.8	0.04	F
FUNDULUS		BALTIMORE	8/21/90	8	2	X			
FUNDULUS	WATER	ELIZABETH	8/21/90	1	1	X	5.1	0.02	M
FUNDULUS	WATER	ELIZABETH	8/21/90	2	1	X	5.8	0.01	F
FUNDULUS	WATER	ELIZABETH	8/21/90	3	1	X	6.7	0.05	F
FUNDULUS	WATER	ELIZABETH	8/21/90	4	2	X	6.6	0.02	F
FUNDULUS	WATER	ELIZABETH	8/21/90	5	2	Х	8.5	0.04	F
FUNDULUS	SEDIMENT	NANSEMOND	10/8/90	R 1	1	X	5.2	0.02	M
		NANSEMOND	10/8/90	2	1	X	5.7	0.04	F
		NANSEMOND	10/8/90	3 🕏	1	Х	5.8	0.05	F
		NANSEMOND	10/8/90	4	1	X	6.5	0.04	М
		NANSEMOND	10/8/90	5	1	Х	8.2	0.06	F
		NANSEMOND:		6	1	X	4.2	0.01	М
	SEDIMENT		10/8/90	7	i ×	X	4.6	0.04	F
	SEDIMENT		10/8/90	8	1	X	3.8	0.02	F
					1	X	4.3	0.04	F
	SEDIMENT		10/8/90	9		X	4.2	0.01	F
FUNDULUS	SEDIMENT	NANSEMOND	10/8/90	10	1	^	7.4	0.01	'

			ANG ANG ANG ANG	580 .	4	×	3.9	0.04	F
FUNDULUS	SEDIMENT		10/8/90	1.1	1		4.2		М
FUNDULUS	SEDIMENT	NANSEMOND	10/8/90	12	1	X	9		F
FUNDULUS	SEDIMENT	DAHLGREN	10/8/90	1	1	X	5.1		F
FUNDULUS	SEDIMENT	DAHLGREN	10/8/90	2	1	X	4.6	•	М
FUNDULUS	SEDIMENT	DAHLGREN	10/8/90	3	1	X	10	• • • •	M
FUNDULUS	SEDIMENT	DAHLGREN	10/8/90	4	1	X		0.05	F
FUNDULUS	SEDIMENT	DAHLGREN	10/8/90	5	1	X	6.7	0.03	F
FUNDULUS	SEDIMENT	DAHLGREN	10/8/90	6	1	X	5.3	0.04	M
	SEDIMENT	DAHLGREN	10/8/90	7	1	X	4.7		F
	SEDIMENT	DAHLGREN	10/8/90	8	1	X	5	0.03	F
	SEDIMENT	DAHLGREN	10/8/90	9	1	X	4.9	0.04	M
	SEDIMENT	DAHLGREN	10/8/90	10.	1	X	6.3	0.03	
FUNDULUS	SEDIMENT	DAHLGREN	10/8/90	11	1	X	4.3	0.02	F
FUNDULUS	SEDIMENT		10/8/90	12	1	X	4.4	0.02	F
			10/8/90	13			22.5		_
FUNDULUS	SEDIMENT		10/8/90	1	1	X	8.7	0.07	F
	SEDIMENT		10/8/90	2	1	X	5.1	0.02	M
	SEDIMENT		10/8/90	3	1	X	5.5	0.01	М
	SEDIMENT		10/8/90	4	1	X	5. 8	0.01	M
	SEDIMENT		10/8/90	5	1	X	7.7	0.06	F
	SEDIMENT		10/8/90	6	1	X	6.8	0.03	M
FUNDULUS	SEDIMENT	INDIAN HD	10/8/90	7	1	X	5. 9	0.04	F
	SEDIMENT			8	1	X	5.2	0.06	F
	SEDIMENT		10/8/90	9	1	X	7.3	0.06	F
	S SEDIMENT		10/8/90		1	X	10	0.08	F
	S SEDIMENT		10/8/90	10	1	x	4.6	<.01	M
	S SEDIMENT		10/8/90		-	x	5.7	0.04	M
	S SEDIMENT		10/8/90		1	x	6.4	0.08	F
	S SEDIMEN		10/8/90		1	X	5.1	0.03	M
	S SEDIMEN		10/8/90		1	X	5	0.04	F
FUNDULU	S SEDIMEN	T POSSUM PT	10/8/90		1		9.5	0.08	F
FUNDULU	S SEDIMEN	T POSSUM PT	10/8/90		1	X	7.6	0.04	M
	IS SEDIMEN		10/8/90		1	Х	3.1	0.02	•••
	S SEDIMEN		10/8/90		1	X		0.02	М
	IS SEDIMEN		10/8/90	8 (1	X	6.5	0.02	F
	IS SEDIMEN		10/8/90	9	1	X	3.3		F
	JS SEDIMEN		10/8/90	10	1	X	4.6	0.03	M
	JS SEDIMEN		10/8/90	11	1	X	4.5	0.01	F
	JS SEDIMEN		10/8/90		1	X	8	80.0	F
EUNDUL	IS SEDIMEN	IT FREESTONE	10/8/9	0 2	1	X	8.1	0.08	
	JS SEDIMEN				1	X	13.3	0.13	F
LONDOF	IS SEDIME	NT FREESTONE			1	X	7.3	0.02	M
LOMDOT:	IS SEDIME	NT FREESTONE			1	X	4.8	0.03	F
FUNDOL	US SEDIMIEI	VT FREESTONE			1	X	4.4	0.03	F
					1	X	7.8		М
	US SEDIMEI			1(0)	1	X	4.7	0.01	М
	US SEDIME				<u>.</u> 1	Х	6.8	0.04	F
FUNDUL	US SEDIME	NT FREESTONE			2	X	6.2	0.03	
		NT FREESTONE			1	X	8.4	0.05	M
	US SEDIME				1	X	12.2	0.28	F
	US SEDIME				1	x	5.3	0.02	M
	US SEDIME					x	5	0.05	
	LUS SEDIME				1	x	7.9	0.17	
	US SEDIME				1	×	6.4	0.06	
FUNDUL	LUS SEDIME	INT MORGANTO	DM 10/8/9	90 5	1	^	E 0.4		

								0.04	14
		MORGANTON	10/8/90	6	1	X	5.3	0.01	M
FUNDULUS	SEDIMENT	MORGANTON	10/8/90	7	1	X	6.5	0.11	F
FUNDULUS	SEDIMENT	MORGANTON	10/8/90	8	1	X	5.8	0.06	F
FUNDULUS	SEDIMENT	MORGANTON	10/8/90	9	1	X	5.5	0.17	F
FUNDULUS	SEDIMENT	MORGANTON	10/8/90	10	1	X	3.2	0.03	F
FUNDULUS	SEDIMENT	MORGANTON	10/8/90	11	2	X	6.7	0.04	M
FUNDULUS	SEDIMENT	CONTROL	9/6/90	1	1	X	5.5	0.04	M
FUNDULUS	SEDIMENT	CONTROL	9/6/90	2	1	X	6.9	0.07	F
FUNDULUS	SEDIMENT	CONTROL	9/6/90	3	1	X	6.3	0.03	М
FUNDULUS	SEDIMENT	CONTROL	9/6/90	4	1	X	8.1	0.08	M
FUNDULUS	SEDIMENT	CONTROL	9/6/90	5	1	X	7.6	0.07	F
FUNDULUS	SEDIMENT	CONTROL	9/6/90	6	1	X	7.2	0.03	F
FUNDULUS	SEDIMENT	CONTROL	9/6/90	7	1	X	7.5	0.06	F
FUNDULUS	SEDIMENT	CONTROL	9/6/90	8	1	X	8.1	0.07	F
FUNDULUS	SEDIMENT	CONTROL	9/6/90	9	1	X	4.7	0.03	F
FUNDULUS	SEDIMENT	CONTROL	9/6/90	10	1	X	6.9	0.44	F
FUNDULUS	SEDIMENT	CONTROL	9/6/90	11	2	X	6	0.04	F
FUNDULUS	SEDIMENT	BALTIMORE	9/6/90	1	1	X	4.6	0.05	F
FUNDULUS	SEDIMENT	BALTIMORE	9/6/90	2	1	X	9.4	0.15	F
FUNDULUS	SEDIMENT	BALTIMORE	9/6/90	3	1	X	8.9	0.15	F
FUNDULUS	SEDIMENT	BALTIMORE	9/6/90	4	1	X	7.8	0.05	F
FUNDULUS	SEDIMENT	BALTIMORE	9/6/90	5	1	X	6.9	0.09	F
FUNDULUS	SEDIMENT	BALTIMORE	9/6/90	6	1	X	9	0.14	F
FUNDULUS	SEDIMENT	BALTIMORE	9/6/90	7	1	X	7.1	0.16	F
FUNDULUS	SEDIMENT	BALTIMORE	9/6/90	8	1	X	3.6	0.01	M
FUNDULUS	SEDIMENT	BALTIMORE	9/6/90	9	1	X	4.6	0.04	F
FUNDULUS	SEDIMENT	BALTIMORE	9/6/90	10	1	X	3.8	0.03	F
FUNDULUS	SEDIMENT	BALTIMORE	9/6/90	11	1	X	4	0.03	M
FUNDULUS	SEDIMENT	WER	9/6/90	1	1	X	6.6	0.09	F
FUNDULUS	SEDIMENT	WER	9/6/90	2	1	X	8.8	0.03	M
FUNDULUS	S SEDIMENT	WER	9/6/90	3	1	X	5.8	0.07	M
FUNDULUS	SEDIMENT	WYER	9/6/90	4	1	X	5.2	0.09	F
FUNDULUS	SEDIMENT	WER	9/6/90	5	1	X	7.4	0.03	F
FUNDULUS	S SEDIMENT	WER	9/6/90	6	1	X	5	0.07	F
FUNDULUS	SEDIMENT	WER	9/6/90	7	1	X	5.7	0.1	F
FUNDULUS	SEDIMENT	WYER	9/6/90	8	1	X	6	0.04	F
FUNDULUS	SEDIMENT	WER	9/6/90	9	1	X	5.7	0.05	F
	SEDIMENT		9/6/90	10	1	X	4	0.01	M
	SEDIMENT		9/6/90	11	1	X	6.1	0.04	F
	S SEDIMENT		9/6/90	12	1	X	5.9	0.13	M
		r ELIZABETH	9/6/90	1	2	X	6.5	0.01	M,F
	•								

Appendix B

Water quality data collected at 1-h intervals with datasonde units during the Potomac River experiments (9/23/90 - 10/2/90). Stations were Dahlgren (DG), Morgantown (MT), Possum Point (PP), Freestone Point (FP) and Indian Head (IH)

	95		
	< **		
.2°			
Ç.		8	

STATION ID : DG

TIME HHMM	TEMP DEG C	PH UNITS	COND MS/CM	SALIN PPT	DO MG/L(PPM)	BATTER/ /OLT3
092390 1400 1500 1600 1700 1800 1900 2000 2100 2200	+21.42 +21.46 +21.46 +21.37 +21.20 +20.95 +20.91 +20.78 +20.70 +20.57	+07.72 +07.73 +07.74 +07.73 +07.91 +07.89 +07.89 +07.84 +07.67	+14.74 +14.86 +14.95 +14.55 +14.26 +14.12 +14.35 +14.33 +14.60 +14.48	+08.33 +08.40 +08.46 +08.27 +08.02 +07.93 +08.06 +08.24 +08.16	+09.08 +08.94 +08.96 +08.84 +09.22 +09.30 +09.34 +08.43 +08.43 +08.23	+05.76 +05.75 +05.75 +05.75 +05.74 +05.74 +05.73 +05.73 +05.73
092490 0000 0100 0200 0300 0400 0500 0600 0700 0800 0900 1000 1100 1200 1300 1400 1500 1400 1500 1500 1600 1700 1800 1900 2000 2100	+20.40 +20.44 +20.23 +20.51 +20.44 +20.32 +20.06 +19.77 +19.59 +19.73 +19.98 +20.11 -20.44 +20.27 +20.32 +20.32 +20.32 +20.33 +20.19 +20.15	+07.57 +07.53 +07.64 +07.41 +07.46 +07.39 +07.41 +07.48 +07.61 +07.66 +07.76 +07.76 +07.76 +07.66 +07.66 +07.68 +07.68 +07.68 +07.68	-14.35 +14.45 +14.45 +14.99 +14.92 +14.83 +14.45 +14.54 +14.57 -14.77 +14.77 +14.77 +14.77 +14.77 +14.77	+08.10 +08.26 +08.31 +08.49 +08.37 +08.37 +08.37 +08.37 +08.33 +0	+08.21 +07.95 +07.98 +06.93 +07.10 +06.81 +07.15 +07.15 +07.15 +08.25 +08.59 +08.59 +08.11 +08.11 +08.12 +08.11 +08.11 +08.11 +08.11 +08.11 +08.15 +08.11 +0	+05.73 +05.73 +05.73 +05.73 +05.73 +05.72 +05.72 +05.72 +05.72 +05.72 +05.72 +05.72 +05.72 +05.72 +05.72 +05.72 +05.72 +05.72 +05.72 +05.72 +05.72 +05.72

TIME	TEMP DEG C	PH	COND MS/CM	SALIN PPT	00 MG/L(PPM)	BATTERY VOLTS
092590 0000 0100 0200 0300 0400 0500 0600 0700 0800 0900 1000 1100 1200 1300 1400 1500 1400 1500 1400 1700 1800 1900 2000 2100 2200 2300	+19.73 +19.68 +19.54 +19.53 +19.60 +19.32 +19.32 +18.82 +18.82 +19.63 +19.37 +19.15 +20.15 +20.15 +20.15 +20.15 +20.15 +20.15	+07.75 +07.74 +07.79 +07.82 +07.57 +07.53 +07.53 +07.47 +07.60 +07.52 +07.61 +07.50 +07.52 +07.64 +07.73 +07.75 +07.83 +07.79 +07.81 +07.75 +07.68	+14.68 +14.55 +14.55 +14.53 +14.68 +14.82 +14.79 +14.73 +14.72 +14.72 +14.72 +14.77 +14.77 +14.79 +14.80 +14.80 +14.80 +14.80 +14.80 +14.80	+08.27 +08.27 +08.21 +08.19 +08.36 +08.32 +08.37 +08.37 +08.35 +08.35 +08.35 +08.35 +08.35 +08.35 +08.37 +08.41 +08.37 +08.37	+08.35 +08.35 +08.36 +08.44 +07.85 +07.55 +07.55 +07.55 +07.55 +07.55 +07.55 +07.55 +07.55 +07.55 +07.55 +08.55 +08.83 +08.83 +08.83 +08.83 +08.83 +08.83 +08.83	-05.71 +05.71 +05.71 +05.71 +05.70 +05.70 +05.69 +05.69 +05.69 +05.69 +05.69 +05.70 +05.70 +05.70 +05.70 +05.70 +05.70 +05.70 +05.70 +05.70 +05.70
092690 0000 1100 0200 0300 0400 0500 0600 0700 0800 0900 1000 1100 1200 1300 1400 1500 1600 1700 1800 1900 2000 2100 2200 2300	+20.06 +19.98 +19.98 +19.73218 +19.199.10 +199.188 +199.155 +199.155 +190.155 +200.00 +200.00 +200.00 +200.00 +200.00	+07.73 +07.71 +07.71 +07.81 +07.89 +07.84 +07.60 +07.57 +07.57 +07.66 +07.70 +07.70 +07.70 +07.93 +08.10 +08.07 +07.98 +07.98 +07.98 +07.98 +07.98 +07.98 +07.98 +07.91 +07.93 +07.93	+14.85 -14.85 +14.85 +14.85 +14.85 +14.80 +14.80 +14.80 +14.83 +14.83 +14.83 +14.83 +14.83 +14.83 +14.87 +16.87 +16.87 +16.87 +17.87 +17.87 +17.87 +17.87 +17.87 +17.87 +17.87 +17.87 +1	+08.37 +08.37 +08.37 +08.37 +08.37 +08.37 +08.37 +08.37 +08.37 +08.37 +08.37 +08.37 +08.37 +08.47 +0	+07.82 +08.50 +08.54 +08.51 +08.23 +09.04 +09.11 +10.00 +09.65 +09.65 +09.65 +09.64	#055.5.5.5.5.5.5.5.5.5.5.5.5.5.5.5.5.5.5

TIME HHMM	TEMP DEG C	PH BH	COND MS/CM	SALIN PPT	DO MG/L(PPM)	BATTERY VOLTS
092790 0100 0100 0200 0300 0400 0500 0600 0700 0800 0800 1100 1200 1400 1400 1400 1500 1400 1700 1800 1700 1800 2000 2100	-19.74 +19.73 +19.58 +19.64 +19.47 +17.43 +19.09 +19.18 +19.30 +18.67 +19.36 +19.36 +19.48 +19.44 +19.44 +19.43 +19.44 +19.54 +21.53 +21.20 +21.25	+07.71 +07.90 +07.92 +08.02 +08.07 +08.05 +08.05 +07.59 +07.59 +07.87 +07.81 +07.75 -07.72 +07.73 +07.73 +07.93 +07.93 +07.94	+14.84 +14.75 +14.63 +14.55 +14.53 +14.53 +14.53 +14.57 +14.89 +14.71 +14.71 +14.74 +14.74 +14.75 +14.75 +14.75 +14.75 +14.75 +14.75	+08.33 +08.33 +08.33 +08.19 +08.19 +08.19 +08.35 +08.35 +08.35 +08.33 +0	+09.44 -09.47 +09.56 +09.85 -10.04 +09.91 +09.91 +09.91 +07.81 -08.87 -08.42 +08.42 +08.42 +08.42 +08.60 -07.89 +07.89 +09.93 +09.93 +09.93 +09.93	+05.68 +05.68 +05.65 +05.65 +05.65 +05.66 +0
2200 2300	+21.25 +21.25	+07.90 +07.81	+14.45 +14.60	+08.14	+09.57 +09.00	+05.68
092890 0000 0100 0200 0300 0400 0500 0500 0700 0800 0900 1000 1200 1300 1400 1500 1400 1500 1400 1500 1400 1500 15	**21.25* +20.75 +20.74 +20.74 +20.53 +20.53 +20.53 +20.57 +20.61 +20.61 +20.61 +20.61 +20.61 +21.33 +20.61 +21.33 +20.51 -21.37 +21.37 +21.37	+07.81 -08.12 +07.79 +07.79 +07.89 +08.15 +07.86 +07.89 +07.89 +07.98 +07.98 +07.98 +07.98 +07.98 +07.98 +07.98 +07.98 +07.98 +08.13 +08.20 +08.23 +08.27 +08.27 +08.27	+14.62 +14.67 +14.72 +14.63 +14.63 +14.55 +14.55 +14.55 +14.53 +14.53 +14.53 +14.55 +14.55 +14.55 +14.55 +14.55 +14.56 +14.66 +16.66 +1	+08.27 +08.30 +08.35 +08.31 +08.26 +08.19 +08.27 +08.27 +08.32 +08.19 +08.27 +08.21 +08.21 +08.21 +08.23 +08.24 +08.24 +08.24 +08.24	+10.70 +10.92	+05.48 +05.47 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.67

TIME HHMM	TEMP DEG C	PH UNITS	COND MS/CM	SALIN PPT	DO MG/L(PPM)	BATTERY VOLTS
092990 0000 0100 0200 0300 0400 0500 0600 0700 0800 0900 1000 1100 1200 1400 1500 1400 1500 1400 1500 1400 1500 1400 1500 1400 1500 1400 1500 1400 1500 1600 1700 1800 1900 2000 2100 2200	+21.42 +20.95 +20.97 +20.97 +21.08 +21.08 +21.08 +21.07 +20.91 +20.91 +21.16 +21.16 +21.16 +21.16 +21.16 +21.54 +21.54 +21.58	+08.17 +08.12 +07.99 +08.07 +07.95 +07.90 +07.75 +07.74 +07.86 +07.99 -07.96 +07.97 +07.97 +07.97 +08.20 +07.94 +08.13 +08.07 +08.01 +08.16	+14.55 +14.57 +14.57 +14.55 +14.55 +14.55 +14.55 +14.55 +14.55 +14.53 +14.53 +14.53 +14.53 +14.55	÷08.25 +08.23 +08.22 +08.22 +08.22 +08.21 +08.21 +08.21 +08.25 +08.25 +08.25 +08.21 +08.15 +08.14 +08.14 +08.14 +08.15 +08.21 +08.21 +08.21 +08.21	-10.71 +10.24 +09.44 +10.03 +09.32 +08.46 +08.46 +08.46 +08.54 +08.54 +09.16 +09.16 +09.10 +09.37 +09.37 +09.37 +09.37 +09.37 +09.37 +09.37 +09.37 +09.37 +09.37	+05.67 +05.67 +05.67 +05.67 +05.67 +05.67 +05.66 +0
2300 093090 0000 0100 0200 0300 0400 0500 0600 0700 0800 0900 1000 1100 1200 1300 1400 1500 1400 1500 1400 1500 1600 1700 1800 1700 2000 2100 2300	+21.50 +21.54 +21.54 +21.54 +21.50 +21.50 +21.50 +21.54 +21.54 +21.54 +21.54 +21.54 +21.54 +21.54 +21.54 +21.54 +21.55 +21.63 +21.63 +21.63 +21.63 +21.63 +21.63 +21.63	+08.03 +08.14 -07.37 +07.79 +07.79 +07.59 +07.59 +07.46 +07.56 +07.56 +07.55 +07.51 +07.52 +07.51 +07.52 +07.52 +07.71 +07.77 +08.02 +07.79 +07.71	+14.56 +14.57 +3.4.92 +14.70 +3.4.4.90 +3.4.4.4.4 +14.70 +3.4.4.4 +14.70 +3.4.4.4 +14.70 +3.4.4 +14.70 +3.4.4 +14.70 +3.4.4 +3.4.4 +3.4.4 +3.4.4 +3.4.4 +3.4.4 +3.4.4 +3.4.4 +3.4.4 +3.4.4 +3.4.4	+08.21 +08.21 +08.22 -08.44 +08.37 +08.51 +08.51 +08.69 +08.67 +08.81 +08.85 +08.85 +08.85 +08.85 +08.65 +08.76 +08.76 +08.64 +08.64 +08.65	+09.84 +09.83 +09.83 +09.83 +09.83 +09.83 +07.84 +08.05 +06.84 +06.82 +07.17 +07.19 +07.19 +07.39 +07.39 +06.91 +06.97 +07.91 +08.64	+05.67 -05.67 -05.66 -05.66 -05.66 -06.66

	TIME	TEMP DEG C	PH UNITS	COND MS/CM	BALIN PPT	CO NG/L(PPM)	BATTERY POLIS
	100190						
	0000	+21.50	F07.74	-39.43	+08.55	-08.67	-05.43
	m100	#21.4Z	£07.84	T. 8. 11	+08.57	-07.90	-05.65
	0200	+21,37	+07.91	-14.91	+09.44	+08.01	+05.63
	0300	+21.50	+07.93	→>>.<<	+08.55	+08.33	+05.53
	0400	+21.54	-07.83	+>>,<<	+08.67	+08.15	-05.53
	0500	+21.54	+07.79	+>>,<<	+08.57	+07.83	+05.63
	0600	+21.53	+07.70	+>>.<<	+08.65	+07.58	+05.63
	0700	+21.46	+07.84	+>>.<<	+08.57	.+07.73	+05.63
	0800	+21.25	+07.92	+15.00	+08.49	+08.04	+05.43
	0900	+21.12	+08.00	+14.93	+08.45	+08.37	÷05.52
	1000	+21.16	+07.95	+>>,<<	+08.52	-08.45	-05.63
	4100	+21.20	-07.Sl	+30.04	+08.51	+07.72	+05.63
	1200	+21,12	467,78	-50 4	-08.67	-07.87	5.51
	1300	±21.08	<i>-</i> 07.70	+\$\$.3€C	+08.73	+07.61	-05.52
	1400	+21.08	÷07.58	+ 50,44	÷08.57	+07.48	+05.52
	1500	421.12	+07.68	+>>,<<	+08.70	+07.29	+05.63
	1600	+21,29	+07.77	+>>,<<	+08.45	+08.32	+05. <u>4</u> 3
	1700	+21.42	+07.90	+>>.<<	+08.57	+07.99	+05.63
	1800	+21,20	+07.71	+>>,<<	+08.74	+07.62	+05.63
	1900	+21.25	+07.75	+>>,<<	+08.73	+07.74	+05.63
	2000	+21.33	+07.77	+>>,<<	+08.71	+07.78	+05.63
	2100	+21.33	+07,71	+>>.<<	+08.71	+07,55	+05.63
	2200	+21.20	+07.87	+>>.<<	+08.70	+08.30	+05.53
	2300	+21.16	+07.96	+>>,<<	+08.58	+08.64	+05.63
	100290		2				
	0000	+21.12	+07.84	+>>,<<	+08.71	+08.27	+05.43
	0100	-20.71	+07.36	±55 , €€	-08,72	-08,38	±05,40
	0200	+20.70	+07.73	+>>,<<	+08.65	-07.21	+05.62
	030¢	+20.78	+07.62	÷>>, <<	÷08.60	+07.27	÷05.62
	9400	+20.74	+07.30	455.KK	+08.57	+07,50	-05.52
	0500	+20.82	÷07.87	->>,<<	+08.51	+08.05	+05.62
	0600	+20.78	+07.90	⊶3¢,বংক	+08.60	-08.19	-05.52
	0700	+20.74	+07.84	+>>,<<	+08.61	+08.20	+05,42
17 post-cal	0800	+15.25	+06.98	+07.23	+03.67	+08,46	+05.60
	0700	÷16.30	+09.95	+30,44	+09,71	÷08.93	-05.59
_ اهه-۱۵مم ۱۵	<u> 1000</u>	+18.71	÷0 7, 70	- >0 , > 4	+09.74	+08.33	4-05.51
	1100	+20,11	+09.74	2 - 1 - 1 - 1	+09.79	+08.07	-05.62
	7.500	-25.70	+08.89	-00.40	+0 0 ,00	+08.85	- +175. 52
	T200	+22.94	+09,14	÷00.34	+00.00		+05.55
	1400	+25.13	+09.23	+00.47		+08.55	÷05.65
	1500	+25.47	+09.22	100.44	+00.00	±07.89	-05.56
	1600	+25.64	.+09-23	+00.48	+00.00	-07.61	+05.57
	£700	+25,47	+09.23	÷00.44	+00.00	107.17	-05.67
	1800	-22.05	±09.22	+00,47	-00.00	±07,50	+05.66
	1300	+24.67	+09.21	÷00.47	+00.00	+07.51	-05.66

B-5

STATION ID : MT

TIME HHMM	TEMP DEG C	PH STINU	COND MS/CM	SALIN PPT	00 MG/L(PPM)	BATTERY WOLTS
092390 1430 1530 1630 1730 1830 1930 2030 2130 2230	+22.34 +22.18 +23.06 +22.26 +22.51 +23.57 +23.44 +22.89 +21.37 +21.37	+07.49 +07.51 +07.45 +07.49 +07.45 +07.30 +07.30 +07.53 +07.53	+13.14 +13.27 +13.54 +13.13 +13.55 +>>.<< +>>.<< -14.39 +13.16 -14.20	+07.31 +07.39 +07.57 +07.57 +07.57 +08.65 +08.82 +08.10 +07.33 -07.99	+07.04 +07.23 +06.88 +07.11 +06.74 +05.50 +05.51 +06.56 +07.26	+05.60 +05.57 +05.59 +05.58 +05.58 +05.58 +05.58 +05.58 +05.58 +05.58
072470 0030 0130 0230 0330 0430 0530 0630 0730 0830 0730 1030 1130 1230 1430 1530 1530 1530 1530 1530 1530 1530 2030 2030 2230	+21.73 +21.29 +21.20 +20.95 +21.12 +20.99 +20.91 +21.50 +20.87 +22.81 +21.45 +20.70 +30.70 +30.70 +20.70 +21.12 +21.29 +21.29 +21.29 +21.29 +21.29 +23.74 +23.74 +23.56	+07.40 +07.43 +07.49 +07.49 +07.47 +07.46 +07.45 +07.34 +07.55 +07.55 +07.55 +07.55 +07.52 +07.52 +07.52 +07.52 +07.52 +07.52 +07.52 +07.52 +07.52 +07.52	+13.73 +13.12 +13.03 +12.03 +12.50 +12.50 +12.50 +13.59 +14.22 +12.73 +14.73 +11.35 +11.35 +11.35 +14.92 +14.92 +14.85 +14.85	+07.68 +07.30 +07.24 +07.15 +06.91 +06.99 +07.28 +06.97 +08.01 +07.79 +06.85 +06.81 +06.85 +06.81 +06.85 +06.81 +06.85 +06.81 +06.85 +06.81 +06.85 +06.83 +08.93 +08.93 +08.93 +08.93	+06.45 +06.88 +06.84 +07.01 +07.01 +07.00 +06.78 +07.41 +06.33 +07.59 +07.59 +08.13 +07.38 +0	+05.57 +05.55 +05.55 +005.55 +005.55 +005.55 +0005 +00

TIME	TEMP DEG C	PH UNITE	COND MS/CM		DO MG/L(PPM)	BATTERY LOUTS
772570 0030 0130 0230 0330 0430 0530 0630 0730 0830 0930 1030 1230 1430 1430 1430 1430 1430 1430 1430 14	+22.74 +22.18 +21.71 +21.50 +21.08 +21.25 +21.25 +23.40 +23.02 +22.60 +22.81 +22.50 +22.50 +22.50 +22.75 +21.75 +21.50 +23.61 +22.72 +23.23 +22.77 +23.23	+07.31 +07.34 +07.36 +07.38 +07.38 +07.40 +07.39 +07.34 +07.24 +07.25 +07.27 -07.31 +07.31 +07.37 -07.46 +07.40 +07.45 +07.40	->>. < < +14.61 +14.64 +13.78 +13.129 +13.228 +>>. < < < +14.30 +13.228 +>>. < < < < < < < < < < < < < < < < < <	+08.83 +08.25 +08.15 +07.72 +07.34 +07.40 +08.80 +08.61 +08.63 +08.63 +08.48 +08.47 +07.14 +07.14 +07.14 +06.73 +08.33 +08.37 +0	+05.38 +06.29 +06.49 +06.49 +06.49 +06.29 +06.20 +05.20 +05.20 +05.20 +05.20 +05.20 +05.20 +05.20 +05.20 +05.42 +05.20 +06.20 +0	-05.54 -05.54 -05.54 -05.54 -05.54 -05.54 -05.54 -05.54 -05.54 -05.54 -05.54 -05.54 -05.54 -05.54 -05.55 -05.54 -05.55
2330 092690 2030 0230 0230 0430 0530 0630 0730 0830 0930 1030 1230 1430 1430 1430 1430 1430 1430 1530 1430 1530 1750 1830 1750 1830	+22.98 +21.80 +21.42 +21.42 +21.50 +21.55 +22.55 +22.56 +22.55	+07.33 +07.35 +07.35 +07.47 +07.46 +07.35 +07.35 +07.37 +07.40 +07.41 +07.41 +07.41 +07.44 +07.50 +07.58 +07.59 +07.29 +07.31 +07.31 +07.31	+>> . < < +>> . < < +14.29 +14.95 +14.97 +14.97 +14.37 +14.35 +14.35 +14.35 +14.35 +14.85	+08.84 +08.63 +08.63 +08.63 +08.04 +08.46 +08.55 +08.55 +08.55 +08.50 +08.51 +08.51 +07.58 +07.58 +07.13 +07.13 +08.75 +08.75 +08.75	+06.02 +06.03 +06.05 +06.05 +06.01 +06.01 +06.05 +06.15 +06.15 +06.41 +06.42 +06.42 +06.35 +06.42 +06.63 +06.77 +07.51 +08.08 +06.05 +06.05	+055.5555555555555555555555555555555555

TIME	TEMP DEG G	DNITS PH	IOND MS/OM	BALIN TEQ	DO (SPM)	BATTERY OLTS
092790 0030 0130 0230 0330 0430 0530 0630 0730 0830 0730 0830 1430 1430 1430 1530 1430 1530 1430 1530 1430 1530 1430 1530 1430 1530 1530 1730 1830 1930 2030 2030	+22.09 +21.01 +21.92 +21.33 +20.95 +20.78 +20.44 +20.44 +20.44 +21.42 +22.22 +23.06 +23.65 +23.65 +23.60 +22.60 +22.01 +22.51 +22.94	+07.34 +07.37 +07.38 +07.45 +07.45 +07.45 +07.45 +07.32 +07.32 +07.27 +07.27 +07.34 +07.35 +07.35 +07.35 +07.35 +07.35 +07.35 +07.35 +07.35	+14.49 +14.49 +13.341 +13.41 +12.41 +12.41 +12.45 +12.45 +12.45 +12.45 +13.45 +14.67 +13.97 +13.97 +13.97 +13.40 +14.07 +13.40 +13.41 +14.07 +13.41 +14.07 +13.41 +14.07 +14.07 +15.00 +14.07 +15.00 +16.00 +	+08.57 +08.17 +08.17 +07.44 +07.48 +06.99 +06.87 +06.87 +06.99 +07.46 +07.98 +07.90 +07.13 +07.53 +07.46 +07.53 +07.48 +07.49 +08.14 +07.48 +07.48 +07.49 +08.99 +08.89 +0	-05.29 +06.50 +06.50 +06.07 +077.00 +0777.00 +0777.00 +0777.00 +0777.00 +05.83 +05.83 -066.13 +066.13 +066.14 +066.14 +066.14 +066.14 +066.14 +066.14 +066.14 +066.14 +066.14 +066.14 +066.14 +066.14 +066.15	+05.52 +05.52 +05.51
2230 2330	+23.44 +23.27	+07.29 +07.29	+>>.<<	+08.73 +08.68	+05.47 +05.72	+05.51 +05.51
092890 0030 0130 0230 0430 0530 0530 0730 0930 1030 1230 1430 1530 1430 1530 1630 1630 1730 1730 1730	+22.77 +22.94 +21.96 +21.97 +21.49 +21.49 +21.49 +21.30.80 +21.30.42 +20.80 +21.30.42 +22.30.42 +23.30.42	+07.29 +07.30 +07.35 +07.35 +07.35 +07.35 +07.35 +07.35 +07.23 +07.23 +07.23 +07.23 +07.33 +07.33 +07.71 +07.72 +07.72 +07.72 +07.72 +07.72 +07.72 +07.72 +07.72	+ · · · · · · · · · · · · · · · · · · ·	+08.80 -08.76 +08.53 +08.55 +08.35 +08.13 +07.62 +07.62 +08.75 -08.95 -08.84 +08.03 -08.84 +08.03 -08.64 +08.03 -08.76 +08.76 +08.76 +08.76 +08.76 +08.76	+05.58 +05.72 +05.72 +06.05 +06.00 +06.25 +0	+05.51 +05.51 +05.551 +05.555 +05.555 +055.555 +055.551 +055.551 +055.555 +

TIME HMMM	TEMP DEG ©	PH UNITS	COND MS/CM	SALIM PPT	DO -G/L(PPM)	BATTERY VOLTS
092990 0030 0130 0230 0330 0430 0530 0430 0730 0830 0930 1030 1130 1230 1430 1530 1430 1630 1730 1830 1930 2030 2130 2230	+22.89 +22.35 +22.72 +23.02 +22.13 +22.05 +21.58 +21.71 +21.29 +21.29 +21.54 +23.44 -22.47 +22.85 +22.81 +22.81 +22.81 +22.81 +22.81 +22.91 +22.91 +22.91 +22.91 +22.91 +22.91 +22.91 +22.91 +22.91 +22.91 +22.91 +22.91	+07.54 -07.58 +07.47 +07.55 +07.37 +07.34 +07.36 +07.40 +07.40 +07.50 -07.55 +07.55 +07.55 +07.59 +07.59 +07.59 +07.59 +07.46 +07.42	+ > > . < < + > > . < < + > > . < < + > > . < < + > > . < < + + > > . < < < + > > . < < + > > . < < + > > . < < + > > . < < + + 14 . 35 5 + 14 . 35 5 + 13 . 43 5 + 13 . 43 5 + 13 . 43 5 + 13 . 43 5 + 13 . 43 5 + 14 . 5 > . < < < < < + > > > . < < < < < < + > > > > > 14 . 5 7 4 + 13 . 44 5 7 1 + 13 . 44 5 7 1 + 13 . 44 5 7 1 + 13 . 44 5 7 1 + 14 . 5 6 6 7 1 + 14 . 5 6 7	+08.74 +08.71 +09.12 +08.81 +08.38 +08.20 +08.07 +07.44 +07.50 +08.51 -08.51 +08.45 +08.45 +08.45 +07.50 +07.50 +07.11 +08.31	-06.89 +06.89 +06.99 +06.05 +06.05 +06.06 +06.26 +06.60 +06.77 -06.77 -06.77 -06.77 +07.21 +07.21 +07.44 +07.44 +07.44 +07.45 +08.16 +06.16 +06.16	+05.51 +05.50 +05.50 +05.50 +05.50 +05.50 +05.49 +05.49 +05.30 +05.49
2330 093090 0030 1130 0230 0330 0430 0530 0430 0730 0830 0930 1030 1230 1430 1530 1430 1530 1630 1730 1830 1930 2030 2030 2030	+23.65 +23.61 -22.72 +22.47 +22.77 +22.13 +22.05 +21.88 +21.75 +21.67 +21.84 +22.78 +22.51 +22.51 +22.51 +22.51 +22.26 +21.75 +21.88 +21.75	+07.37 +07.37 -07.53 +07.51 +07.51 +07.56 +07.46 +07.45 +07.45 +07.45 +07.45 +07.54 +07.54 +07.54 +07.59 +07.59 +07.61 +07.67 -07.58	+14.71 +>> +>> +>> +>> +>> +	+08.61 +08.72 +09.39 +09.14 +08.73 +08.17 +08.17 +08.17 +07.23 +07.52 +08.64 +09.35 +09.35 +09.35 +08.44 +07.85 +07.85 +07.63	+05.45 +06.45 +06.45 +06.57 +06.75 +06.77 +06.73 +06.73 +06.73 +06.57 +06.57 +06.57 +06.57 +06.57 +07.73 +0	+05.50 +05.49 +0

TIME HHMM	FEMP DEG C	STINU	COND MS/CM	SALIN PPT	QO MG/L(2PM)	BATTERY VOLTS
100190						
0030	+22.30	F07.52	+13.31	÷07.42	+07.30	-05.43
0130	+24.84	+07.26	+>>,<<	+09.32	+05.20	-05.48
0230	+24.15	+07.23	+>>,<<	+08.34	-05.35	+05.47
0330	+23.61	+07.32	+>>.<<	+08.59	+05.71	-05.49
0430	+23.10	T07.31	ナリン。くぐ	+08.80	-05.52	+05.47
0530	+22.72	+07.36	514.89	-08.4T	405.73	-05.48
9630	+21.63	+07.41	+13.65	±07.53	-06.47	+05.48
0730	+21.50	<u>-07.43</u>	+13.84	+07.75	+06.58	+05.48
0830	+21.54	+07,40	+13.69	+07.56	-06.54	+05.48
0930	+21.13	-07.34	+13.92	+07.81	+06.15	+05.47
1030	+21.20	+07.42	+13.11	+07.29	+06.17	+05.47
1170	+21.19	+07.57	-12.88	-07.15	-07.23	+05.47
1270	-21.FI	-07.52	-13.32	+07.43	-07,48	-05,47
1370	+23.53	-07.28	+ 50 . 44	+08.95	+05.49	±05.48
1450	+25.54	÷07.25	+>>.<<	-09,24	+05.36	+05.48
1530	+24.25	+07.31	- 12.44	+09.12	+05.71	+05.49 +05.49
1630	+24.63	+07.37	+>> . < <	+08.92	+05.94	+05.47
1730	+24.46	+07.36	+>>.<<	+09.09	+05.73 +05.66	+05.48
1830	+23.78	+07.33	+>>.<<	+09.19	+08.15	+05.48
1930	+22.18	+07.80	+13.66	+07.64 +07.51	+08.80	+05.48
2030	+22.09	+07.99 +07.34	+13.45 +13.14	+07.32	+08.44	+05.48
2130 2230	+21.71 +21.50	+07.59	+12.90	+07.16	+07.90	+05.47
2330	+23.95	+07.41	+>>.<<	+08.59	+06.75	+05.47
2000	. 23. 73	107.41				
100290						
0030	+24.75	+07.31	+ 23 . 3 4	+09.16	+05.91	+05.48
1100	-24.41	±07.54	State of the state	-08.99	-OE. ?2	-05.48
0230	+23.40	-07.25	+0,	+09.35	-05.18	-05.48
0330	+23.48	+07.20	->>. <	+10.17	+04.51	+05.43
0430	+22.26	+07.25	F>>. < <	+10.50	-04.39	+ 5.48
A530	+22.85	± 07.19	+>>=<<	+10.34	+04.07	±05.47
0630	+22.22	H07.27	+22.55	+09.75	-04.75	-05.47
0730	+21.71	+07.33	+>9,<<	+09.10	+05.46	+05.47
0830	+16.73	+06.87	+07.04	+03.56	+07.86	+05.46
್ರಿ9ಸ್ಟರ	+19.30	+06.86	F00.46	+00.00	+08.74	+05.46
1030	+20.44	±97.20	-00.37	±+00.00	+07.78	-05.47
1170	+20.49	⊧97.21	-00.37	F00.00	-07.54	+05.47
1230	+21.50	<u>-07.23</u>	+00.40	+00.00	±07.30	+05.47
1330	+24.71	+07.21	+00.41	+00.00	-07.42	+05.48
1450	+25.59	F07.20	F00.41	+00.00	-07.00	+05.48
1530	+25.54	+07.17	-00.41	+00.00	-05.70	÷05.48
1430	+25.14	.HO7.15	+00.41	+00.00	+04.51	+05.48

STATION ID : PP

TIME	TEMP	=4	COMD	BALIN	50	SATTERY
HMM	DEG C	STINU	MS/CM	PPT	MG/L(PPM)	VOLTS
092390						
-1100	+23.32	-07.71	+90.62	00.00	+34.40	-05.67
1200	+21.58	-07.57	+00.33	+00.00	+07.97	+05.69
1300	+21.63	-07.48	+00.31	+00.00	±07.78	+05.68
1400	+21.54	+07.45	+00.31	+00.00	+07.70	+05.68
1500	+21.63	+07.44	F00.31	+00.00	+07.64	÷05.58
1600	+21.67	+07.47	+00.31	+00.00	+07.74	+05.67
1700	+21.63	+07.45	÷00.31	+00.00	+07.58	+05.67
1800	+21.46	+07.39	+00.31	+00.00	+07.47	+05.57
1900	+21.37	-07.37	+00.29	#00.00	+07.42	+05.67
2200	+ 21.57	-47.55	-00.35	>0,66	-08.33	⊣್ತ್.≃೧
E100	+21.37	-07.42	-00.31	+00.00	-07.58	+05.55
2200	+21.37	-07.42	+00.35	+00.00	-07.50	+05.05
2300	+21.29	-07.44	+00.36	+00.00	+07.65	405.66
092490						_
0000	+21.12	+07.37	-00.32	+00.00	+07.52	+05.66
0100	+20.95	+07.41	+00.32	+00.00	+07.78	+05.65
0200	+20.78	÷07.33	+00.30	+00.00	+07.44	+05.45
0300	+20.65	+07.36	+00.30	+00.00	+07.32	+05.45
0400	+20.49	+07.36	+00.30	+00.00	+07.31	+05.65
0500	+20.35	+07.35	+00.32	+00.00	+07.32	+05.35
0500	+20.40	+07.34	+00.30	+00.00	+Q7.33	+05.65
0700	+20.57	+07.J1	+00.30	+00.00	+07.27	+05.64
0800	+20.70	+07.33	+00.32	+00.00	÷07.43	+05.64
1200	-Z0.74	5307 DE I	HOO.UZ	+00.00	407.JB	+05.44
F556	-20.74	-07.35	÷00,32	+00.00	+07.42	+05.64
7.700	÷20.82	407.36	+00.J0	÷00.00	+07.36	+05,54
1200	420.91	-07,36	÷00.32	+00.00	-07,50	+05.54
1300	÷20,95	-Q7,35	+00.JZ	+00.00	+07.56	+05.64
1400	420.75	-07.Ja	-00.32	-00.00	+07.36	F05.54
1500	+20.37	-07.37	+00.50	+00.00	+07.58	÷05.64
1600	+20.95	-07.36	+00.32	+90.00	+07.42	+05.64
:300	+20,91	+07.37	+00.32	+00.00	+07.29	÷05.64
<u> </u>	÷20.87	407.38	+00.32	+00.00	+07.44	+05.63
1500	+20.91	-07.35	**+00.30	+00.00	+07.58	-05.63
2000	+20.87	+07.32	+00.32	+00.00	+07.35	+05.53
2100	+20.82	407.3Z	+00.32	+00.00	÷07,69	+05.80
2200	+20,78	-07.45	°+80.34	+00.00	-07 . 94	+05.53
Z300	-20.SZ	+07,44	-00.34	-00.00	+07.07	+05.63
		207				

TIME 	TEMP DEG 🕠	PH UNITS	COND MS/CM	EALIN OPT	00 1671.(FPM)	BATTERY VOLTS
172570				A. (A. 1975)	+07.73	-05.53
9600	-20.74	-07,46	+00.34	-00.00	+07,75	-05.63
0100	÷20.70	+07.45	-00.34	-00.00	-07.51	+05.63
0200	+20.53	+07,43	+00.34	+00.00	+07.55	+05.62
0300	+20.32	+07.38	+00.32	+00.00	+07.26	+05.62
0400	+20.32	+07.33	+00.32 +00.30	+00.00	+07.14	+05.62
0500	+20.19	+07.33 +07.31	+00.30	+00.00	+06.97	+05.62
0600	+20.19 +20.19	+07.31	+00.30	+00.00	+07.21	+05.62
0700	+20.17	+07.32	+00.30	+00.00	+07.31	+05.62
0800 0900	+20.40	+07.33	+00.32	400.00	+07.45	+05.62
1000	+20.44	-07.35	+00.32	+00.00	+07.47	+05.62
190	+20.57	-07.J5	+00.30	-00.00	+07.41	+05.62
1200	-27.51	+07.36	-00.32	-00.00	+07.50	-35.52
1300	+20.70	-07.36	-00.32	-00.00	+07.50	-05.52
1400	+20.82	407.36	+00.32	-00.00	+07.59	÷05.52
1500	+20.95	-07.39	+00.32	400.00	+07.82	÷05.⊍Z
1400	+21.03	+07.42	+00.32	÷00.00	+07.95	-05.62
1700	+20.99	+07.42	+00.32	+00.00	+07.93	+05.62
1800	+20.95	+07.43	+00.32	+00.00	+07.97	+05.62
1900	+20.95	+07.42	+00.34	+00.00	+07.92	+05.42
2000	+20.82	+07.37	+00.32	+00.00	+07.62	+05.61
2100	+20.78	+07.35	+00.32	+00.00	+07.65	+05.61
2200	+20.78	+07.41	+00.32	+00.00	+08.01	+05.51
2300	+20.82	±07.43	±00.34	+00.00	+08.00	+05.61
092590					26.31	
0000	+20.78	+07.44	+00.34	+00.00	+08.01	+05.01
OO IN	-22.65	-07.45	#90.34	###O \$22	-08.17	-415,61 05 -1
0200	-20.51	-()7.44	+00.54	+00.60	+08.05	-05.61 -05.61
9300	+20.49	F07.43	+00.34	±00.00	+08.01	#05.51
0400	+20.55	+07.43	+00.52	+00.00	+08.04	+05.61
9500	#20.2J	+07.43	+00.32	÷00.00	+08.05	+05.50
D600	+20.15	-07.43	+00.32	+00.00	+08.95 +07.90	+05.60
0700	+20.06	+07.42	+00.32	+00.00	+07.75	+05.50
0800	+20.15	+07.42	+00.32	+00.00	+07.90	+05.50
0900	+20.27	+07.40	+00.32	+00.00	-07.98	+05.50
1000	+20.32	+07.40	+00.32	+00.00	+08.01	+05.60
1100	+20.40	+07.41	** +00.32	+00.00	-08.01	-05.50
1200	+20.49	+07.42	+00.32	+00.00	+07.95	+05.50
1300	+20.53	+07.42	+00.32	-00.00		+05.50
1400	+20.40	-07.43	+00.32	+00.00	3.7	+05.60
1500	+20.57	+07.45	+00.34	+00.00		+05.50
1600	+20.14	+07.46	+00.32 +00.32	+00.00		+05.60
1700	+20.49	F07.45	+00.32	-00.00	3.5 5.4	+05.60
1800	+20.44	-07.47	+00.32	+00.00		+05.60
1900	+20.35	+07.47	+00.32	+00.00		+05.60
2000	+20.40	+07.44	+00.32	+00.00		+05.50
2100	+20.44 +20.44	+07.41 +97.58	-00.32		5 7 71	§05.60
2200 2700		-07.48	+00.32	-00.00		+05.50
2700	+20.55	-9/.40	F (, 10) •	.5.5 . 50		

TIME	TEMP DEG 0	PH UNITE	DOND MS/CM	BALIN	00 4371(29M)	SATTERY STJQ.
092790 0000 0100 0200 0300	+20.40 +20.36 +20.32 +20.19	+07.43 +07.43 +07.41 +07.44	+00.34 +00.32 +00.32	+00.00 +00.00 +00.00 +00.00	-07.86 +07.87 +07.76 +08.01	-05.59 +05.59 +05.59 +05.59
0400 0500 0400 0700 0800 0900	+20.06 +20.02 +19.89 +19.77 +19.73 +19.68	+07.42 +07.40 +07.39 +07.42 +07.43	+00.32 +00.32 +00.32 +00.32 +00.32 +00.33	+00.00 +00.00 +00.00 +00.00 +00.00	+07.73 +07.76 +07.62 -07.63 +07.47 +07.48 +07.43	+05.59 +05.59 +05.59 +05.59 +05.59 +05.59 +05.59
1000 1100 1200 1300 1400 1500	+19.98 +20.23 +20.40 +20.53 +20.19 +20.19	+07.36 +07.52 +07.51 -07.32 +07.33 +07.33	+00.32 +00.32 -00.32 +00.32 +00.32 +00.32	+00.00 +00.00 -00.00 +00.00 +00.00	+07.52 -07.33 +07.57 +07.48 -07.50	+05.59 +05.59 +05.59 +05.59
1500 1700 1800 1900 2000 2100	+20.65 +21.29 +21.75 +21.58 +21.46 +20.78	+07.32 +07.46 +07.49 +07.48 +07.44 +07.36	+00.32 +00.31 +00.31 +00.31 +00.31 +00.32	+00.00 +00.00 +00.00 +00.00 +00.00	+07.39 +08.00 +08.12 +08.02 +07.87 +07.48	+05.37 +05.39 +05.59 +05.59 +05.59
2200 2300 092890 0000	+20.44 +20.40	+07.33 +07.35 +07.38	+00.32 +00.32 +00.32	+00.00	+07.37 +07.55 +07.55	+05.58 +05.59 -05.58
7144 0206 0300 0400 0500	+20.44 +20.49 +20.44 -20.36 +20.40	-07.41 -07.41 +07.41 +07.40 +07.35	+00.74 +00.74 +00.74 +00.72 +00.72	+00.00 +00.00 +00.00 +00.00 +00.00	-07.68 -07.84 +07.71 +07.73 +07.48	-05.58 +05.58 +05.58 +05.56 +05.58
0605 0700 0800 0900 1000	+20.23 +20.23 +20.15 +20.23 +20.23	+07.33 +07.30 +07.29 +07.27 +07.25	+00.32 +00.32 +00.32 +00.72 +00.32	-00,00 +00.00 +00.00 +00.00	+07.07 +07.08 +07.00 +07.08 +07.03	+05.58 +05.58 +05.58 +05.58 +05.58
1100 1200 1300 1400 1500 1400	+20.40 +20.49 +20.40 +20.44 +20.49 +20.57	+07.27 +07.31 -07.34 +07.35 +07.36 #07.35	+00.02 +00.34 +00.34 +00.05 +00.43 +00.34	+00.00 +00.00 +00.00 +00.00 +00.00 +00.00	+07.21 +07.46 +07.53 +07.61 +07.67 +07.61	+05.58 +05.58 +05.58 -05.57 +05.37 +05.57
1700 1800 1900 2000 2100	+21.03 +20.74 +20.61 +20.57 -30.83	+07.34 +07.32 +07.32 +07.31 +07.30	+00.32 +00.32 +00.32 +00.32	+00.00 +00.00 +00.00 +00.00 +00.00	+07.73 +07.57 +07.48 +07.51 +07.45 -07.44	+05.87 +05.87 +05.37 +05.37 +05.87
2300 221	-10.37 -20.74	+07.30 +07.30	-00.32 -00.32	-00.00 -00.00	-07.54	4.5.57

TIME	TEMP DEG C	PH UNITS	COND MS/CM	SALIM PPT	90 MG/L(PPM)	BATTERY OLTS
 (92990						_
0000	-20.61	+07.51	-00.38	+00.00	-yr.59	-05.57
1100	+20.65	+07.37	-00.33	+00.00	+07.72	-,35.37
0200	+20.65	+07.37	+00.45	+00,00	+07,85	+05.57
0300	+20.51	+07.38	+00.47	+00.00	+07.70	+05.57
0400	+20.El	÷07.40	-00.45	+00.00	-07.90	+05.57 ∸05.57
0500	+20.49	÷07.36	-00.33	+00.00	-07.77	+05.57
0600	+20.40	+07.35	+00.34	+00.00	+07.74	+05.57
0700	+20.40	+07.33	+00.34	+00.00	+07.65	+05.57
0800	÷20.32	+07.33	+00.32	+00.00	+07.52 +07.55	-05.36
0700	+20.32	+07.33	+00.32	+00.00		+05.54
1000	+20.55	+07.52	+00.32	+00.00	-07.44 -07.52	-05.56
1100	-20.01	+07.71	-00.02	-90.00	9007.5D	8.89
1550	유민이 . 왕 고	-9€,5€		400.00	-07.88	-05.36
1300	+20.78	+07.31	-00.34	400.00	+07.68	-05.76
<u>400</u>	+20.74	÷07.55	400.38	+00.00	-07.60	-05.55
1590	-20.57	+07.33	-00.43	-00.00	+07.59	-05.35
1600	+20.61	+07.52	+00.42	+00.00	+07.36	-05.54
1700	+20.74	+07,31	+00.40	+00.00	+07,72	-05.55
1800	+20.99	+07.33	+00.36	+00.00	+07.59	+05.55
1900	+20.82	+07.53	+00.38	+00.00	+07.56	+05.54
2000	+20.74	+07.32	+00.38	+00.00	+07.53	+05.54
2100	+20.70	÷07.32	+00.34	+00.00	+07.45	+05.56
2200	+20.74	F07.31	+00.58	+00.00	+07.44	+05.56
2300	+20.65	±07.30	÷00.34	+00.00		
MPIG90				100.00	-07:34	+05.55
9990	-20.61	+07.27	-00.32	+00.00 +30.00	7.15	ed 5 •d° 5
5100	-20.51	FG7.27	-00.38	+00,00	-07.77	-05.56
0200	+20.u5	FQ7.74	+00.45	+00.00	+07.83	495.86
9300	+20.70	+07.05	-00.49 =	+00.00	-07,79	-05,58
Q400	-20,65	-07.56	+00.59 +00.57	+00.00	+07.50	+05.56
0500	+20.61	+07.36	-00.37	±00.00	-07,81	+05.56
->60°-	+20.a1	£07.35	+00.43	+00.00	+07.74	+05.56
5700	+20.49	+07.35 +07.30	+00.45	+00.00	+07.47	-05.56
0800	+20.44	+07.29	-00.32	+00.00	+07.37	F05.55
0900	+20.05	-07.27	-do.54	+00.00	-07.55	#N/5.55
<u>1</u> 000	+20.44 +20.57	+07,27	8-80.I4	-00.00	æ07.∑	
1100	+20.37	+07.27	-50.72	400.00	-07.50	-05.35
1200	-20.32	+07.30	-00.04	-00.C0	∓07.50	-05.55
1100	+20.95	+07,20	+()0.4()	±00.€0	-07.56	a)5.05
1400	+20.95	+07.35	+00.55	+00.00	-07.77	-03.55
1500	+20.79	+07.36	+00.63	+00.00	+07.70	⊧05.35
1.400	+20.77	+07.37	-00.65	-00.00	; +07.65	445.55 ·
1700	+21.03	-07.36	+00.63	+00.00	÷07.87	-05.35
1800	+21.03	+07.35	-00.48	±00.00		⊕05.55
1900	+20,59	+07.35	+00.44	+00.00		-05.55
2000	+20.77	+07.03	-00.40	+00.90	-07.79	-95.05
2100	-20.77	-e7T	400.JS	g +00.04	22 10 2	-115.1E
1200 1200	-20.75	-07.46	-00.35	+00.0		an6.55

HHMM	TEMP DEG C	SUMITS	COND	SALIN Spt	93 MG/1.(PPM:	SATTERY JULIS
100190						
0000	+20.87	+07.57	+00.36	-00.00	+07.65	+05.55
0100	+20.82	+07.32	-00.36	-00.00	+07.52	-u5.55
0200	+20.74	+07,30	-00.40	+00.00	+07.49	÷05.55
0300	+20.78	+07.36	+00.49	+00.00	÷07.77	+05.55
0400	+20.74	+07.05	±00.53	+00.00	+07.58	-05.53
2500	+20.70	÷07.36	-00.76	+00.00	+07.75	-05.85
6400	+20.65	+07.37	+00.74	+00.00	+07.79	+05.55
0700	+20.57	-07.36	÷00.49	÷00.00	+07.63	-05.55
0800	+20.40	607.US	-00.41	#00.00	+07.50	+05.55
0900	+20.36	-07.51	+00.36	÷00.00	+07.41	-05.54
1000	+20.32	÷07.30	+00.34	+00.00	+07.47	+05.54
1100	+30.00	-4 - 7 - 1 T	-24).34	÷∪0.0€	-07.48	-05.54
11000	+20°.85	-117,33	- 10,54	10 .JJ	=7.637 . 3. 4	-65.00
TIOO	+20.71	+07.05	-00.34	+00.00	+07.76	-05.54
1400	+21.20	-07.32	-00.I4	+00.00	+-17.03	FU5.54
1500	F21.42	+97.55	-00.TE	-00.00	-07.54	+05.54
1500	+21.25	-07.39	+00.42	+00.00	+08.05	+05.33
1700	+20.99	+07.36	-00.40	+00.00	+07.59	÷05.34
1800	+21.08	-07.53	+00.48	+00.00	+07.77	+05.54
1900	+21.03	+07.35	+00.44	+00.00	+07.85	÷05.54
2000	+21.03	+07.36	+00.42	-00.00	+07.72	-05.54
2100	+21.16	÷07.37	+00.33	+00.00	+07.76	÷05.54
2200	+20.91	+07.33	+00.56	+00.00	+07.71	+05.54
2300	+20.74	±07.32	+00.34	+00.00	+07.5a	+05.54
100290						
9000	+20.57	+07.12	೬೧೧, 75	-0.00 - 0.1N	407:45	-05.54
	750 . T.			- 6	7.42	#15,54
2200	420 . 7·3	±07.ĪΞ	+00.49	≈00.00	÷07.57	+05.54
0300	+20.78	+07.36	+00.55	+00.00	∸ 07.79	+05.54
9490	+20.78	+07.35	+00.78	+00.00	+07.81	+05.54
0500	+20.74	+07.38	±00.95	+00.00	+07.83	÷05.54
3600	*20.78	m±07.54	+01.10	+00.09	+07.82	-05.54
0700	+20.61	-07.36	+00.89	+00.00	+07.77	+05.54
0800	+20.44	÷07.35	+00.55	+00.00	+07.73	+05.54
) 9 00	4E0.E3	±07.54	+00.47	±00,00	+07,61	+05.55
1.000	-20.12	-07.57	-00,37	400 .(0)	<u> </u>	water.
1100	+17.85	-12.57	Part and	-179	+95.43	:5 , 17

اله م عدر 10 h ح

STATION TO : FR

TIME	-Eb	FH	DOND	SAL IM	23	BATTERY
THIS IN	TEE	-MI-2	HSACM	723	MG/L PPM)	VOLTS
092390						
1030	+20.44	-07.40	+00.29	+00.00	+08.10	+05.84
1130	+20.57	-07.26	+00.29	+00.00	+07.98	+05.83
1230	+20.74	+07.34	+00.29	+00.00	+08.18	+05.83
1330	+20.87	+07.66	+00.27	+00.00	-09,14	-05.82
1430	+20.99	+07.79	+00.31	+00.00	+09.74	+05.82
1530	+20.99	+07.76	+00.31	+00.00	+09.77	+05.82
1630	+21.08	+07.88	+00.31	+00.00	÷09.83	+05.82
1730	+21.20	-09.01	+00.28	+00.00	÷10.06	+05.82
1830	+21.08	+07,86	+00.28	+00.00	+09.85	+05.52
1930	+20,87	-07.63	±00.23	+00.00	+09.50	+05.81
2030	- 2 5.7.4	-07,-1	-00,25	-00.00	-1.8.2	-05.51
2130	+20.57	+07,47	+00.27	+00.00	-05.30	-05.81
ZZ30	+20.47	+07.57	-00.27	+00.00	-08,45	+05.30
2530	*20.36	-07.70	+00.27	+00.00	-MP. 758	-05.80
2000		77 A 7 =		0000		
092470						
0030	+20.19	+07.43	+00.29	+00.00	±09.15	+05.80
0130	+19.98	⊦07.30	+00.29	+00.00	+08.39	+05.80
0230	+19.77	F07.34	+00.29	+00.00	+08.77	+05.80
0330	-19.56	+07.36	±00.29	+00.00	+08.56	+05.79
0430	+19.18	+07.34	+00.30	+00.00	+08.46	+05.79
9530	+17.13	A07.35	+00,30	+00.00	+08.32	+05.79
0630	+19,18	-07,44	-00.30	+00.00	+08. 4 7	+05.78
0730	e+13.92	±07.00	+00.30	+00.00	+08.75	+05.78
0830	+18.88	+07,52	+00.23	-00,80	-08.42	+05.78
1.370	+19.84	a [1평 .]스	-90.IB	- YC . JW	-05	- 15.78
1030	+18.88	-08.2J	+00.28	+00.00	+08.51	+05.78
1130	+18.97	-08.10	-00.28	+00.00	+08.57	+05.78
1230	+19.22	±07.85	+00.30	+00.00	F09.15	-05.77
1330	+19.68	+07.51	+00.29	+00.00	+09.22	+05.77
1430	+20.02	-07.38	+00.29	+00.00	t09.70	-05.77
1530	+20.19	+07.88	÷00.J1	+00.00	+09.96	+05.78
1630	+20.23	F08.03	+00.29	+00.00	+10.05	+05.77
1730	+20.49	+07.75	F00.29	+00.00	+09.78	-05.77
1930	+20.20	÷08.03	H0Q.29	+00.00	+09.57	:05.77
1930	+20.06	:07.72	-00.29	+00.00	+09.31	-05.77
2010	+20.02	+08.77	+00.27	+00.00	+09.J3	+05.77
2130	-19.56	#UB.57	F00.27	-00.00	+10.45	-05.77
1230	+19.55	-48.73	-00.27	-00.00	-09.58	+05.75
2330	-19.47	+08.58	-00.27	-JU. WO	-09.52	-05.76
					25	

FIME	TEMP DEG 0	оч LNITS	COND MS/CM	SALIM	00 ::G/L(2PM:	PATTERY VOLTS
092590						
0030	+19.±0	408.39	+00.27	400.90	-1)9.Js	-05.75
0130	+19.43	-08.32	+00.27	+00.00	-09.59	+05.76
0220	+19.35	-07,40	+00.30	÷00.00	-08.41	+05.76
0330	+18.97	+07.40	+00.28	+00.00	÷0 8. 58	+05.76
0430	+18.75	+07.40	+00.28	+00.00	+08.69	-05.75
2530	+18.59	+07.46	÷00.30	+00.00	+0 8. 53	+05.75
9630	+18.50	+07.53	H00.28	+00.00	+08.53	+05.75
0730	+18.75	+08.13	+00.28	+00.00	+08.95	+05.75
0830	+18.59	F07.27	+00.28	+00.00	+08.20	+05.75
0930	+18.63	+07.Z0	+00.28	+00.00	+07.98	÷05.75
7020	+18.75	+07.66	+00.28	+00.00	+08.01	+05.75
a + A 9	-19.75	-08,22	400,IB	-00.00	+08.67	+05,75
	¥18,92	一/号. /3	0.13	-40,00	-09,11	995.25 -05.75
4.3/5/0	+19.43	-07.71	+00.27	+00.00	+09.12 +08.94	+05.75
1430	+19.77	407.50	-00.27	+00.00	+09.06	+05.75
1530	+20.15	-07.52	+00.27	+00,00	+09.47	-05.75
1530	+20.47	+07.70	-00.29	-00.00 +00.00	+09.34	+05.75
1730	+20.65 +20.44	407.73	+00.29 +00.29	+00.00	+09.17	÷05.75
1830		+07.54	+00.27	+00.00	+08.29	÷05.75
1930	+20.35 +20.44	+07.57 +07.54	+00.29	+00.00	+08.10	÷05.75
Z030 Z130	+20.49	+08.49	+00.27	+00.00	+10.63	+05.75
2230	+20.49	+07.42	+00.29	+00.00	+08.71	+05.75
2230	+20.47	+08.43	+00.27	+00.00	+09.79	+05.75
2000	* ALC: # AL /	1.70 (7,00.127	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		
092690	ee 20 11	27 74		100 00	+08.93	÷05.74
0020	#20.11	H07.74	+00.29	+00.00	-08.20	#05.74
1120	#19.31	-07,54	-00.Z7	+40.40	+08.16	-05.74
0278	+19.77	+07.29	+00.27	+00.00 -00.00	÷08.08	-05.74
0330	+19.51	+07.27	+00.27	+00.00	÷08,06	+05.74
0430	-19.35	-07,21	-00.27	+00.00	+08.10	+05.74
0570 5775	+18.22 +19.13	-07.26 -07.05	+00,28 +00,30	+00.00	+08.32	405,NJ
0430 0730	+19.26	+07.25	+00.30	+00.00	+08.02	+05.73
0830	+19.26	+07.17	+00.32	+00.00	+07.76	+05.73
9730	=12.05	+07.25	+00.30	+00.00	+07,61	+05.73
£030	+19.01	+67,27	+00.30	+00.00	+07,65	405.77
1135	+19,13	-07.49		+00.00	+07.94	+05.77
1230	-19.26	-07,46	-00,30	+00.00	-08.38	-05.77
1330	+19,47	+07.26	÷00.29	-00.00	-08.07	-05.FT
1477	+19,77	-07.71	-00.31	+00.50	+08.42	#05.77
1630	+20.06	÷07.35	+00.31	+00,00	-09.03	÷05.75
1430	-19.74	+07.59	±00.31	400.00	+09.02	+05.73
1770	+20.02	+07.73	#40.E1	+00.00	-09.27	05.73
1930	+19.94	+07.91	.00.51	-00.00	-ე9,93	+05,77
F644	-19.70	F08.15	LOO.II	-NO.74	-10.36	*05.77
2070	±10.74	+07.44	400.71	+00.00	* +08.9I	-05.77
	- 18 - 17	=1075	1 V/1 & T. (.₩C. 1	-08.65	-AE,UX
	1	7	- 10.77		-08.04	-05.57
	-19.64	- -97.54	+00.Iª	-00.00	+08.22	`-OE.TT

FIME	TEMP DEG C	PH UNITS	COND MS/CM	SALIN _PPT	DO MG/L(PPM)	EATTERY VOLTS
092750	÷19.60	+07.38	+00.27	+0 0. 00	-08.4¢	-65.72
0130	419.51	-07,50	+00.32	+00.00	-08.63	-05.23
0230	+19.51	+07.21	+00.29	+00.00	÷08.25	+05.72 +05.72
0330 0430	+19.22 +18.97	+07.29 +07.47	+00.30 +00.30	+00.00	+08.37 +08.93	+05.72
0530	+18.88	+07,48	+00.32	+00.00	-09.04	+05.72
0630	+18.71	+07.23	+00.32	+00.00	+08.56	+05.72
0730	+18.59	+07.22	+00.34	+00.00	+08.44	+05.72
0830	+18.53	-07.25	÷00.32	+00.00	+08.31	+05.72
0930	+18.71	+07.36	+00.32	+00.00	÷08.64	+05.71
1030 1130	÷19.01 ÷19.22	+07.39	+00.30	+00.00	+08.62 +08.86	+05.71 +08.71
1230	717.44 717.43	+07.58 =38.00	+00.33 0.75	+00.00 +30.00	-08.83	50T. F5
1730	+19.45	+07.83	+00.30	+00.00	+08.87	-M3.71
1430	+19.39	-07.77	+00.32	+00.00	+09.15	#05.81
7220	+19.56	-07.35	+00.32	400.00	-10.02	-: [a = 7];
1630	+19.77	÷07.95	+00.31	+00.00	+10.12	-05.71
1730	+20.53	+08.10	+00.31	+00.00	+10.34	aπ95.71
1830	+21.03	+08.13	+00.33	+00.00	+10.95	+05.72
1930 2030	+21.12	+08.11	+00.33	+00.00	+10.87 +10.59	+05.72 +05.72
2130	+21.12 +20.32	+08.01 + 07. 90	+00.33 +00.31	+00.00	+09.78	+05.72
2230	+20.32	+07.43	+00.31	+00.00	+08.65	+05.71
2330	+20.27	+07.35	+00.29	+00.00	+08.14	-05.71
092890						
9939	+R0,36	e07.31	+00.22	+00.00	+07.84	-05.71
177	≈16. <i>1</i> 7	H107,15	400.ZF	-00,00	+07.74	-45.71
0230	-39,II	+07.60	+00.31	+00.00	+08.58	+05.71
0330	-19.98	+07.41	+00.31	+00.00	+0 8. 53	-05.71
0430	+19.98	-07.41	-00.31	+00.00	+08.55	+05.F1
0530	+19.98	+07.56	+00.31	+00.00	+09.02 +09.14	+05.71 +05.71
9430 9730	+20.15	-07.55 +07.58	+00.31 +00.33	+00.00	+09.44	+05.71
0830	+20.02	+07.53	+00.31	+00.00	+08.95	+05.71
0930	+20.02	+07.69	+00.31	+00.00	+09.21	+05.71
1030	+19.74	+07.41	F00.31	+00.00	-08.60	+05.71
1130	+20.15	F07.37	20 +00.31	+00.00	+08.49	+05.71
1230	+20.34	+07.27	+00.31	+00.00	+08.12	+05.70
1330	±20.49	+07.24	+00.31	+00.00	+07.35 +07.97	-05.70 -05.70
1430 1530	*20.40 +20.57	+07.27 +07.53	+00.31 +00.31	+00.00 +00.00	±09.14	+05.71
1630	+20.87	+07.65	+00.31	+00.00	+09.22	+05.71
1730	±21.12	+07.91	+00.30	+00.00	-10.03	+05.70
1830	+21.03	+08.04	+00.30	+00.00	+10.52	÷05.71
1930	+21.46	+08.12	+00.J0	+00.00	+10.67	+05.71
2030	+21.46	+08.04	+00.30	+00.00	+10.42	+05.71
2130	+21.42	+08.13	+00.30	+00.00	+10.60	+05.71
2330 2330	+20.95	-07.79 -07.71	+00.31		-49.53 +09.53	203.71 405.70
-2 201	+20.95	-07.71	+00.31	-00.00	TU7.00	

TIME	TEMP DEG U	PH DNI TS	JOND MS/CM	SALIM PPT	EO MG/LyPPMY	
:92990 :030	+20.87	+07.25	+00.31	-00.00	-09.58	+05.70
0130	+20.75	+07.43	+00.31	+00.00	+08.71	+05.70
0230	+20.61	+07.24	+00.31	+00.00	+07.91	+05.70
0330	+20.70	+07.33	+00.31	+00.00	+08.09 -09.15	+05.70 +05.70
0430 0530	+20.70 +20.70	+07.51 +07.54	+00.31 +00.31	+00.00	±08.50	-05.70
0630	+20.55	+07.39	+00.31	+00.00	+08.59	-05.70
0730	+20.61	+07.62	+00.31	+00.00	-09.10	+05.70
0830	+20.65	+07.61	-00.D1	+00.00	409.28	÷05.70
Q930	+20.57	+07.51	-00.51	-00,00	-09.17	- 3.59
1030	-20,40	-07.38	+00.31	+00.00	-08.85	+05.70
1130	*20.70	-07.53	-00.T1	+00.00	-78.58 -78.57	-1.0.57
1270 1370	-20,73 +20.91	-07,99 -07,51	##0,51 +00.31	=60© -000	-07.95	-v5.54
1430	+20.75 +20.79	-07.Ji	+00.31	+00.00	+07.82	-05.70
1530	+21.03	+07.29	+00.31	-00.00	+07.75	-05.59
1430	+21.12	+07.55	+00.30	+00.00	+08.48	+05.70
1730	+21.25	+07.40	+00.30	+00.00	+08.45	+05.70
1830	+21.42	+07.36	+00.30	+00.00	+08.50	+05.49
1930	+21.46	+07.44	+00.30	+00.00	+09.11 +09.75	+05.69 +05.69
2030	+21.58	÷07.73	+00.32	+00.00	+09.48	+05.69
2130 2230	+21.58 +21.50	+07.45	+00.32 +00.32	+00.00	+09.56	+05.70
2330	+21.20	+07.74	+00.30	+00.00	+08.57	+05.69
093090						
0030	H20.99	+07.28	+00.31	+00.00	-08.13	+05.69
0130	F20.71	+07.76	-00 -51	-14 .00	18.43	- 5.59
0230	F20.87	+07.71	+00.31	+00.90	-08.52	+05.67
0330	+20.74	+07.94	+00.31	+00.00	-08.53	+05.69
0430	+20.82	+07.52	400.51	+00, QQ	+08.51	+05.69 +05.69
0500	+20.87	-07.57	+00.31	+00.00	+08.61 +07.9I	-115.69
0430 0730	+20.37 +20.78	+07.19 +07.19	+00.31 +00.27	+00.00	÷07.90	+05.59
0830	+20.87	+07.28	+00.31	+00.00	F08.21	+05.69
0930	+20.95	+07.22	+00.33	±00.00	-08.18	+05.69
1010	+20.95	F07.25	400.JJ	+00.00	+08.40	-U5.67
1130	+20.95	-07.46	§ +00.J1	-00.00	-08.55	F05.69
1230	+21.20	-07.25	+00.J0	-00.00	+08.10	+05.69 +05.69
1370	=21.25	+07.44	-00.30	+00.90	+07.65 +07.53	-05.57
7220 7430	-21.42 *21.42	⊬07.30 ∸07.45	+00.39 +50.38	+00.00 +00.00	-07.85	⊕05. 5 7
1630	+21.46	VFQ7.37	+30.30 +30.30	+00.00	+07,17	+05.69
1730	#21.75	-07.52	-00.32	-00.00	-08.04	-05.67
7820	-21.67	+07.45	-00.30	F00.00	÷08.09	-05.67
1930	#21.63	+07.22	+00.30	+00.00	+08.02	-05.57
2030	+21.71	+07.34	+00.30	+00.00	- +08.44	+05.67
1130	-31.53	607.JZ	+00.32	+00.00	+08.57 +08.53	+95.69 +05.69
2270	+21.65	-07.30	-00.32	-00.00 -00.00	-08.73	405.68
2330	+21.42	+07.35	-00.02	-UNJ . U	.,0.	. 5 . 4 6

TIME HAMM	TEMP DEG C	PH UNITS	COND MS/CM	SALIN PPT	00 MG/L(FRM)	SATTERY VOLTS
100190						
0030	F21.25	+07.42	-00.30	-00.00	-07.81	+05.68
OFIC	+21.16	÷07.13	-00.30	400,00	+07.54	÷05.68
9230	+20.99	+07.15	+00.51	±00.00	+07,43	+05.68
0330	+20.78	+07.59	+00.31	+00.00	+07.79	+05.58
0430	+20.82	+07.21	+00.31	÷00.00	+07.09	-05.58
0530	+20.70	+07.26	+00.31	+00.00	+07.53	+05.48
0630	+20.53	+07.35	+00.31	+00.00	+07.77	+05.58
0730	+20.32	+07.12	+00.31	+00.00	+07.50	+05.68
0 8 30	+20.15	+07.17	+00.31	+00.00	+07.72	+05.57
୨30	+20.19	+07.22	+00.31	+00.00	+07.85	÷05.57
1030	+20.49	+07.01	+00.39	+00.00	+07.50	+05.67
1170	+20.57	-07,09	-00.37	+00.00	÷07.59	÷05.5√
_130	+30.57	-07.JE	np.37	-440.00	+x,470 , \$150	කැලී.කැදී
1330	+21.03	+07.34	-00.31	+00.00	+07.43	+05.88
1430	+24.33	+07.27	+00.30	-00.00	+07,99	+05.67
1530	+21.42	+07.38	+00.30	+00.00	+07.17	±05.68
1630	+21.29	+07.83	+00.30	+00.00	+07.72	∸05.58
1730	+21.15	+07.50	+00.30	÷00.00	+07.71	+05.67
1930	+21.20	+07.51	+00.30	+00.00	÷07.63	+05.67
1930	+21.33	+07.37	+00.30	+00.00	+08.10	+05.67
2030	+21.25	+07.49	+00.30	+00.00	+08.48	+05.67
2130	+21.03	+07.42	+00.31	+00.00	+08.52	+05.67
2230	+21.03	+07.43	+00.33	+00.00	+08.81	+05.67
2330	+20.99	+07.41	+00.23	÷00.00	∸08.⊖3	÷05.67
100290						
0030	+Z0.74	÷07.J7	+00.31	-00.00	+08,16	+05,47
0130	-20.49	-07.22	-90.51	-00.00	-07.30	-05.a7
0230	+20.19	+07.34	+00.31	+00.00	407.48	÷05.97
<u>0330</u>	+20.05	±07.32	+00.31	400.00	-07,迂止	+05.67
0430	+19.89	407.53	÷00.29	+00.00	+07.52	405,67
0530	+20.11	+07.32	+00.29	+00.00	+07.35	+05.56
0630	+19.87	407.28	+00.29	+00.00	+07.40	÷05.68
0730	+19.85	+07.26	+00.29	+00.00	+07.70	+05.56
0830	+19.77	+07.29	+00.31	+00.00	+07.95	+05.66
0970	+[0.81	F07.75	+00.31	+00.00	+08.25	+05.56
1350	117-46	_±47,~1		-+00.4il		<u> 66</u>
1130	+10.72	+07,00	≈ ±07.40	-03.77	-08.57	#05.66
1230	-20.44	-L0.05	+00.KK	-08.67	+08.47	-05.57

47 post-cal

B-20

STATION ID : I4

TIME	TEMP	24	EOND	EALIN	90	BATTERY
무단인인	DEG Ĉ	LAHTS	MSACM	₽PT	467F(BBW)	VOLTS
292370						
1000	+21,29	±Ω7,24	+00.31	<u>+00.00</u>	+07,68	+05.73
1100	+21.33	+07.09	+00.31	-00.00	+06.76	+05.72
1790	+21.54	÷07.09	+00.31	-00.00	+06.97	+05.71
1300	+21.57	± 07.19	+00.31	-00.00	+07,04	+05.71
1400	+21.42	+07.08	+00.31	+00.00	+06.59	+05.71
1500	+21.46	+07.08	+00.31	÷00.00	+06.95	+05.71
1600	-21.50	*07.10	+00.31	+00.00	±07,09	-05.71
1700	+21.54	+07.13	÷00.33	+00.00	+07,37	+05.71
1800	±21.42	÷07.12	+00.31	+00.00	+07.25	+05.71
_ POC	#21.25	-07.10	400.JJ	-50.00	+07,18	+05.70
2007	<u>≈34.10</u>	* 7 . <u></u> .	-00.51			= 15 mg
2100	-21.15	907.1T	400.J1	-00.00	-1.7.2	-05. (アの
2200	+21.00	-07,14	-00.51	-00.00	+077.4	÷05.70
2700	+21.03	⊫07.13	+00.31	+00.00	-i177	-05.70
092490						
0000	+20.91	+07.15	+00.31	+00.00	+07.40	+05.70
0100	+20.65	+07.16	+00.31	+00.00	+07.50	+05.69
0200	+20.82	+07.12	+00.31	+00.00	407.Ji	+05.69
0300	+20.45	+07.11	+00.31	+00.00	+07.28	+05.69
0400	+20.61	+07,10	+00.31	+00.00	-07.25	+05.69
0500	+20.51	+07.10	+00.31	+00.00	+07.25	+05.57
0400	+20.51	÷07.12	+00.31	+00,00	+07,37	÷05.69
0700	+20.15	-07.12	+00.32	+00.00	+07.02	+05.49
0800	+20.27	+07.10	÷00.31	+00.00	+07,25	+05.69
1990	+20.27	第四页,11	450.51	-00,00	-07.23	-65.67
1000	+20.27	+07.12	-00.31	+00.00	+07.Ja	-05.59
1100	+20.36	+07,13	+00.31	+00.00	+07,42	+05.69
1200	+20.53	+07.1J	+00.31	+00.00	÷07.53	+05.58
1300	+20.70	+07.13	+00.33	+00.00	+07,53	+05.58
1400	+20.82	÷67.14	-00.31	-00.00	÷07.65	÷05.58
1500	+20.91	+07.13	+00.31	+00.00	+07.44	+05.68
1600	+20.70	÷07.10	+00.31	-00.00	+07.33	-05.53
1700	+20.70	-07.09	+00.31	-00.00	+07.27	+05.67
1800	+20.32	F07.10	+00.31	+00.00	+07.17	+05.57
1900	+20.91	-07.12	≈ +00.JI	-00.00	F07.55	-05.67
2000	120.78	+07.10	+00.31	+00.00	+07.47	+05.67
2100	+20.70	+07.12	-00.31	+00.00	+07.19	+05.57
2200	+20.65	+07.11	+00.31	+00.10	-07.43	+05.47
2300	+20.61	+07.13	+00.31	+00.00	-07.51	+05.67
2000	. 20.01		11.0.51	00.00		

TIME	TEMP DEG C	=H UNITS	COND MS/CM	SALIN	00 BA 36/L(PPM)	ATTERY VOLTS
		Shrie				
192590						
2000	+20.56	+07.16	+00.31	+00.00	-07.51	+05.67
0100	+20.36	+07.15	+00.31	+00.00	+07.57	+05.67
0200	+20.40	-07.14	+00.31	+00.00	+07.55 +07.30	+05.67 +05.67
0300 0400	+20.53	+07.09	+00.33	+00.00	+07.30	+05.67
0400 0500	+20.44 +20.36	-07.09 -07.10	+00.33 +00.31	+00.00 +00.00	-07.23	+05,67
0600	+20.20	-07.10	+00.32	+00.00	+07.18	+05.47
0700	+20.15	+07.10	+00.32	+00.00	+07.20	+05.67
0800	+20.15	+07.09	+00.32	+00.00	+07.20	+05.67
0700	+20.15	+07.08	+00.32	+00.00	+07.17	+05.67
1000	+20.15	-07.10	+00.32	-00.00	407.24	+05.67
1100	+20.23	-07.⊃P	+00.CZ	#00.90	+07.E2	-05.66
. I Day	- 217. 27	-57.14	-cQ.71	- () () () ()	-07.27	-05.65
7200	-20.27	-07.14	+00.31	-00.00	+07.55	+05.55
1400	+20.53	-07.14	+00.33	+00.00	+07.60	-05.66
1500	-20.70	#97.12	+00.21	+00.00	-07.53	+05,aa
1500	+20.44	+07.08	+00.31	+00.00	+07.25 +07.25	+05.55 +05.56
1700	+20.44	+07.08	+00.33	+00.00 +00.00	+07.25	+05.66
1900	+20.44 +20.53	+07.07	+00.33 +00.31	+00.00	+07.38	+05.66
1900 2000	+20.33	+07.10 +07.10	+00.33	+00.00	+07.29	+05.56
2100	+20.44	+07.10	+00.33	+00.00	+07.40	+05.56
2200	+20.40	+07.09	+00.31	+00.00	+07.33	+05.45
2300	+20.44	+07.10	+00.33	+00.00	+07.40	+05.65
092694	,					
0000	+20.35	+07.10	E00.31	-00.00	407.T4	-05.65
1403	-20.27	=17.1	-30.01	-30.19	-177.76	-05.55
1200	-20.27	+07.12	+00.01	+00.00	-07.44	+05.65
1300	+20.23	+07.09	+00.32	+00.00	+07.37	+05.65
9400	+20.23	+07.98	+00.32	+00.00	+07.34	+05.45
2500	-20.19	-07.09	-00.32	+00.00	+07.15	+05.45
0400	+20.05	-07.08	-00.32	+00.00	+07,26	+05.65
0720	±20.02	+07.08	+00.32	+00.00	+07.23 +07.18	#05.65 +05.65
0800	+20.06	+07,07	+00.32	+00.00	+07.27	-05.65
0700	+30,02	-07.08	+00.32	+00.00	-07.51	-05.65
1000 1100	+20.02 +20.02	+07,09 -07,09	-00.32 -00.32	+00.00	-07.35	-05.65
1200	#30.02	+07,08	+00.34	+00.00	+07.50	+05.65
1200	+20.17	+07.10	+00.32	+00.00	+07.46	-05.e5
_A00	-20.20	407.11	+00.33	=00.00	+07.45	-05.64
1500	+20.19	=07,14	+00.32	+00,00	+07,69	∸05.65
4400	+20.36	-07.12	+00.33	+00.00	+07.54	-05.04
1,700	+2Q.27	+07.08	-00.33	+00.00	÷07.36	-05.64
1800	+20.23	=07.07	+00.53	-00.00	-07.54	+05.84
TAGO	-IO.E3	#D7,03	+00.33	÷00.00	+07,22	+05.64
2000	+20.27	+07.04	+00.31	+00.00	+07.33	+05.64
2766	-20,27	t07.55	+00.51	-00.00	+07.31 =87.33	+05.64
1221	-20,19	707.12	-10.12	-00.50	-07.52	-05.64
2500	+20.1T	-47.15	FD9.JJ	-90.00	-177 - 2-	17 J 1 U 7

TIYE	TEMP	7:0	COND	EALIM	00	BATTERY
-HMM	CEG C	CNITE	MS/ICM	PPT	*G/L[PPM)	- MOLIS
072790						
0000	±20.15	+07.12	+00.JZ	±00.00	-07.47	+05.54
0100	+20.19	+07.09	+00.32	400.00	+07,42	+05.54
0200	+20.15	+07.12	+00.34	+00.00	+07.43	+05.64
0300	+20.05	+07.12	+00.32	+00.00	+07.56	+05.63
Q40Q	+20.02	+07.22	+00.32	+00.00	+07.61	+05.54
0500	+20.02	-07.06	-00.32	+00.00	+07.46	+05.54
0600	+20.02	+07.06	+00.32	+00.00	+07.35	+05.63
0700	+19.94	+07.0-6	+00.32	+00.00	+07.25	+05.4J
0900	÷19.89	+07.06	+00.32	+00.00	+07.25	+05.43
9 9 00	+19.81	+07.07	+00.02	+00.00	+07,32	+05.63
1000	+19,29	±07.10	+00.32	+00.00	+07.45	+05.62
- F O C	+20,62	-07, <u>1</u> 1	-00.5I	-00.30	÷3/7,43	
- Z/5 O	-25.J≙	96 FO.	-00	See Control	#1702 p 3 1	
_ 50 O	+20,99	-07.19	+00.71	÷00.50	-17.50	d Min konsil
1400	521.19	- 07,21	-00.51	Fig. 7.	408.01	-05.50
1300	#21.50	⊝연필·교실	+00.II	+00.00	408.13	-05.63
1500	+20.70	+07.LD	+00.33	+00.00	+07.79	+05.63
1700	+20.23	+07.11	+00.32	+00.00	+07.52	÷05.52
1900	+20.15	+07.08	+00.32	+00.00	+07.36	+05.52 +05.62
1900	+20.06	+07.09	+00.32	+00.00	+07.41	+05.62
2000	+20.11	+07.08	+00.32 +00.32	+00.00	+07.22	+05.62
2100 2200	+20.06 +20.27	+07.07 +07.08	+00.32	+00.00	+07.17	+05.62
2300	+20.27	+07.11	+00.31	+00.00	+07.51	+05.62
2000	20:4-	10/12	100.01	100100	0 · • • ±	32.02
092890						
0000	+20.27	+07.19	+00.51	+90.00	⊶07,65	-05.62
E 1 1/25/25	*20.25	51/8 · 22	-02.72			
9200	#20.0I	+0,7.22	400.TZ	+00.810	+08.01	+05.62
-1200	+20.02	-07.20	+00.32	+00.00	+07.92	+05.62
94.00	± 20.19	÷07.11	+00.02	-00.00	+07.63	<u>+05.62</u>
0500	+20.35	+07.17	+00.J1	+00.00	+07.95	+05.62
5600	+20.19	+07.10	-50.02	+00.00	+07.72	-05.52
0700	+20.23	+07.10	+00.32	+00.00	+07.58	+05.52
0800	+20.11	+07.10	+00.02	+00.00	+07.40 +07.41	+05.52 +05.62
0500	-20.06	+07.69	+00.04	+00.00	+07.25	+05.62
1000	+20.11	+07.08	+00.32	+00.00	+07.42	-05.62
1100	+20.17	+07.11	₹ +00.32	+00.50 +00.00	+07.59	+05.62
1200	+20.27	+07.12	+00.31 +00.31	+00.00	+07.90	-05.62
1300 1400	+20.53 +20.74	+07.13 +07.23	+00.33	+00.00	+08.34	-05.62
1500	+22.05	+07.32	+00.30	+00.00	+08.75	-05.62
1600	+21.75	g+07.33	+00.30	+00.00	-08.87	+05.62
1700	+20.70	+07.15	-00.19	+00.00	+08.15	+05.62
1800	+20.27	+07.08	-00.31	+00.00	+07.51	+05.62
1900	+20.32	H07.11	+00.33	+00.00	+07.56	+05.62
2000	+20.23	+07.07	+00.32	+00.00	+07.41	+05.61
2100	+20.15	+07.06	+00.32	+00.00	±07.32	+05.51
IZOO	+20.36	+07.08		-00.00	+ 17,45	<u> 5</u> 05.51
2300	+20.27	+07.08	+00.31	+00,00	-O7:48	405.51

TIME HHMM	TEMP DEG C	PH STING	COND MS/CM	SALIM PPT	20 MG/L(PPM)	BATTERY UCLIS
200000						
092990 0000	+20.49	-07.13	700 71		-07.53	+05.51
0100	+20.49	+07.11	+00.31 +00.31	+00.00 +00.00	-07.65	+05.61
9200	+20.57	+07.14	+00.31	+00.00	+07.74	+05.61
0300	+20.49	+07.16	+00.31	+00.00	+07.88	+05.61
0400	+20.49	+07.15	+00.31	+00.00	+07.84	-05.51
0500	+20.44	+07.12	+00.31	+00.00	+07.70	+05.51
0600	+20.32	+07.11	. +00.33	+00.00	+07.62	+05.61
0700	+20.32	+07.09	+00.33	+00.00	+07.69	+05.61
0800	+20.32	+07.08	+00,31	+00.00	+07.54	+05.51
0900	+20.27	+07.08	+00.33	+00.00	+07.59	H05.51
1000	+20.23	+07.08	+00.32	+00.00	+07.52	+05.61
1100	+20.JZ	+07.09	+00,51	+00,00	+07,43	405,61
1200	-20,40	+:177		-110 . O.P	F07,43	-05.51
1300	+20.91	+07.18	÷00.31	+00.00	HO8.14	+05.50
1400	+20.91	+07.17	+00.33	+00.00	+08.23	-05.61
1500	+20.95	-07.17	+00.31	+00.00	+08.17	-05.5I
1600	+20.82	+07.15	+00.33	+00.00	+08.05	+05.61
1700	+20.99	+07.19	+00.33	+00.00	+08.34	+05.61
1800	+21.12	+07.20	+00.33	+00.00	+08.42	+05.61
1900	+20.78	+07.13	+00.31	+00.00	+07.84	+05.61
2000	+20.53	+07.09	+00.31	+00.00	+07.72	+05.50
2100	+20.44	+07.07	+00.31	+00.00	+07.55	+05.60
2200	+20.53	+07.08	+00.31	+00.00	+07.57	+05.60
2300	+20.49	+07.07	+00.31	+00.00	+07.58	+05.60
007000						
093090	+20,53	. 07 . 50	.00 71	.00.00	107 10	105 40
0000 0110	-20.57	-07.07 -07.09	+00.31 +00.51	+00.00	407.40 -07.73	+05,60 -05,60
0200	+20.65	+07.11	+00.31	-90.00 +00.00	+07.76	-05.60
0300	+20.73	+07.15	+00.29	+00.00	+07.95	+05.60
0400	+20.78	+07.17	+00.51	+00.00	+07.99	÷05.60
0500	+20.74	+07.22	+00.51	+00.00	+08.26	+05.60
0600	+20.74	+07.14	+00.31	+00.00	F07.59	+05.50
0700	+20.49	+07.08	+00.29	+00.00	+07.65	+05.60
0800	+20.49	+07.09	+00.31	+00.00	+07.59	+05.60
0900	+20.44	+07.09	+00.31	+00.00	+07.70	+05.50
1000	+20.36	+07.08	+00.31	+00.00	+07.55	+05.30
1100	+20.49		** +00.31	+00.00	+07,82	405.8Q
1200	+20.61	+07.10	+00.31	400.00	+07.73	+05.60
1390	+E0.95	-07.15	o+00.33	+00.00	+08.17	+05.50
1400	+21.15	+07.18	+00.33	+00.00	+08.34	+05.40
1500	+21.58	-07.17	+00.31	±00.00	+08.15	+05.50
1600	+E1.03	+07,17	÷00.31	+00.00	+08.11	405.60
<u> </u>	±21.08	+07.19	+00.29	+00.00	+08.15	+05.50
1900	+21.08	±07.13	+00.31	÷00.00	+08.17	+05,60
1900	+20.82	+07.12	+00.33	+00.00	÷08.05	-05.50
2000	+20.82	+07.13	+00.33	+00.00	+08.02	+05.60
2100	+20.78	+07.14	+00.33	+00.00	+08.04	+05.59
2200	+20.78	-Q7,11	-00.JJ	+00.00	+08.03	-05.59
2700	+20.TB	+07.14	+00.33	-00.00	+08.Ga	+05.59

	TIME HHMM	TEMP DEG C	PH UNITS	COND MS/CM	SALIN PPT	00 MG/L(2PM)	BATTERY VOLTS
	100190						
	0000	+20.74	+07.14	+00.23	±00.00	-08.08	+05.59
	0100	+20.61	-07.13	+00.31	+00.00	-08.03	+05.59
	0200	+20.51	+07.10	+00.33	+00.00	÷08.00	+05.59
	0300	+20.53	+07.12	-00.31	+00.00	+07.98	+05.59
	0400 0500	+20.49	+07.12 +07.11	+00.29 +00.29	+00.00 +00.00	-08.03 +07.95	+05.59 +05.59
	0600	+20.36 +20.32	+07.12	+00.31	+00.00	+07.75	+05.59
	0700	+20.27	+07.13	+00.31	+00.00	+07.97	+05.59
	0800	+20.36	÷07.10	+00.31	+00.00	+07.95	+05.59
	0900	+20.36	+07.09	+00.31	+00.00	+07.83	+05.57
	1000	+20.53	÷07.05	+00.31	+00.00	+07.75	+05.59
	1100	+20.57	-07,08 -07,865	+00.33	-00.00 -00.00	-07.71	-05.57
	2200	-20.55	#.17 133		400.T)	-47.35	A 37
	1300	+20.53	=97,58	+00.11	-00,00	-07,ĒĪ	-15.39
	1400	-70,81	907.47	-00.13	400.0C	+07.77	+05.39
	30	-20.78	E07.1.	+00.53	+00.00	+07,99	-05.59
	1400	+21.29	+07,17	+00.31	+00.00	+08.38	+05.59
	1700	+21.20	+07.34	+00.29	+00.00	+08.84	+05.59
	1800	+21.08	+07.28	+00.31	+00.00	+08.87	+05.59
	1700	+20.78	+07.19	+00.31	+00.00	+08.40	+05.59
	2000	+20.91	+07.19	+00.31	+00.00	+08.33	+05.58
	2100	+20.57	+07.06	+00.33	+00.00	+07.74	+05.58
	2200	+20.65	+07.07	+00.33	+00.00	+07.72	+05.58
	2300	+20.70	+07.09	+00.33	+00.00	+08.01	+05.58
	100290						
	0000	+20.53	+07.09	+00.33	+00.00	+08.02	+05.58
	2100	425.23	=07.09	-00.53	-00.00	0-08.WE	E05.79
	0200	+20.56	+07.09	+00.31	+00.00	+07.91	+05.58
	0300	+20.23	+07.07	+00.32	+00.00	+07.75	-05.58
	0400	+20.36	+07.17	+00.31	+00.00	+08.21	F05.58
	0500	+20.27	+07.18	+00.30	+00.00	+08.27	+05.53
	0600	+20.19	+07.17	+00.30	+00.00	+08.25	+05.57
	0700	+20.11	+07.18	+00.30	+00.00	+08.23	+05.57
	008C	+20.23	+07.11	+00.32	+00.00	+08.05	+05.57
	0900	+20.27	+07,10	+00.71	+00.00	+08.01	+05.57
11 7 post- cal	1000	+15.84	+07.00	+02.41	+00.84	+10.08	+05.58
1	1100	+19.09	-06.91	+03.25	+01.30	+09.46	+05.58
10 post-cal	1200	+22.98	÷10.00	+>>,=(<	+08.89	+08.57	+05.57
y	1100	+25.47	÷09.79	+>>. <<+	+09.05	+08.63	+05.40
			20				

	p#		
	e e	25	
	u ^c		

Appendix C

Water quality conditions reported in test containers during all water column tests

ge se

140

j.

Table C-1: Experiments conducted with Fundulus spp. (Fs), Palaemonetes pugio (Pp), Cyprinodon variegatus (Cv) and Eurytemora affinis (Ea) using water samples collected from the Wye R. (WR), Patapsco R. (PR), and the Elizabeth R. (ER) from 8/14/90 to 8/21/90

Date	Test Species	Station	Temp (C)	Sal (ppt)	DO (mg/L)	рН
8/14/	======== 00			=======		
0/14/	Fs	WR	26.60	15.5	7.3	7.75
	13	WE	26.20	16.0	7.2	7.79
		PR	25.30	14.5	7.3	7.77
			25.50	15.5	7.5	7.74
		ER	25.30	20.0	7.3	7.61
			25.30	20.0	7.2	7.61
		CONTROL	25.60	15.5	7.1	7.94
			26.30	15.5	7.4	7.95
	Pp	WR	24.60	15.5	4.3	7.51
	•		24.60	15.5	5.3	7.59
		PR	25.40	16.0	6.5	7.68
			24.20	16.0	5.1	7.46
		ER	24.50	20.0	2.7	7.08
			24.80	20.0	2.6	7.12
		CONTROL	24.70	15.5	5.7	8.10
			24.50	16.0	5.8	8.01
	Cv	WR	25.20	15.5	3.8	7.85
			24.90	15.0	3.7	7.86
		PR	25.10	16.0	7.2	7.74
			25.00	16.0	7.4	7.78
		ER	25.20	20.0	6.4	7.48
			25.20	20.0	6.5	7.50
		CONTROL	25.20	15.5	3.8	7.85
			25.10	15. 5	3.7	7.86
	Ea	WR	25.30	16.0	5.2	7.80
			25.50	16.5	5.0	7.75
		PR	25.20	16.5	5.3	7.70
			25.20	16.5	5.5	7.74
		ER	25.50	*	5.0	7.56
			25.70	20.0	4.9	7.50
		CONTROL	25.30	16.0	4.5	7.96
. 0/15	/ 0.0		25.20	16.0	4.4	7.95
8/15/				75.0	6 6	*
	Fs	WR	27.00	15.0	6.6 6.8	*
		PR	2.6.20	15.0	7.4	*
		ER	26.00	20.0	6.9	*
	Dw	CONTROL	27.00	15.0 15.0	2.6	*
	Pp	WR	25.20	15.0	3.9	*
		PR	24.60 26.00	15.0	5.0	*
		FR	25.00	15.0	3.2	*
		ER	25.20	20.0	*	*
		Ľ.K	25.20	20.0	*	*
		CONTROL	25.20	15.0	6.6	*
		CONTROL	25.10	15.0	6.1	*
			23.10	10.0	J	

	CV	WR	25.20	15.0	6.50	* }
	•	PR	25.50	15.0	6.1	*
		ER	25.30	20.0	6.5	*
			25.30	20.0	5.6	*
		CONTROL	25.30	15.0	2.1	*
	Ea	WR	25.20	15.0	_	*
		PR	25.80	15.0	6.5	*
		ER	25.50	20.0	4.3	*
		CONTROL	26.00	16.0	5.0	*
8/16/90						_
	Fs	WR	24.00	15.0	5.6	*
		PR	25.50	15.0	6.8	*
		ER	25.30	20.0	6.6	*
	_	CONTROL	25.20	15.0	6.2	*
	Pp	WR	25.50	15.5	4.2	*
			25.00	16.0	4.2	*
		PR	25.20	15.0	*	*
		ED	25.00	15.5	*	*
		ER	26.00	21.0	*	*
		CONTROL	25.20	20.5 16.0	5.3	*
		CONTROL	25.20 25.70	15.5	5.8	*
	Cv	WR	25.70	15.5	4.8	*
	CV	PR	25.50	15.0	6.0	*
		ER	25.50	20.0	2.7	*
		CONTROL	25.20	15.0	3.4	*
	Ea	WR	25.00	15.0	6.2	*
	Lu	PR	25.70	15.0	7.7	*
		ER	25.50	20.0	4.3	*
		CONTROL	25.20	15.0	5.4	*
8/17/90						
, ,	Fs	WR	26.80	14.5	3.7	7.30
		PR	25.90	13.5	7.0	7.68
		ER	26.00	20.0	6.9	7.60
		CONTROL	26.40	15.0	7.0	8.11
	Рp	WR	25.70	15.0	6.0	7.76
	_		25.30	15.0	6.3	7.78
		PR	25.00	13.5	4 . 5	7.51
			25.20	13.5	*	7.29
		ER	24.90	21.0	5.7	7.55
			25.10	20.5	5.1	7.45
		CONTROL	24.90	14.5	6.5	8.20
			25.50	15.0	6.5	8.14
	CV	WR	25.30	15.0	3.0	7.47
		PR	25.70	14.5	6.4	7.86
		ER	25.50	20.5	2.2	7.17
	_	CONTROL	25.20	15.0	4.4	7.91
	Ea	WR	25.50	15.5	6.5	7.86
		PR	25.50	15.0	4.0	7.53 7.30
		ER	25.00	20.5	4.8	7.30
0 /10 /00		CONTROL	25.00	15.0	5.4	1.33
8/18/90	□ ~	7.775	26.10	15 0	6.3	*
	Fs	WR	26.10	15.0	6.6	*
		PR	25.90	14.0	0.0	••

		ER	26.00	20.0	6.5	*
		CONTROL	26.10	15.0	7.2	*
	Pp	WR	25.20	15.5	5.8	*
	-		25.10	16.0	5.2	*
		PR	25.70	13.5	5.4	*
			25.20	13.5	4.9	*
		ER	24.90	20.0	6.3	*
			25.00	20.0	6.1	*
		CONTROL	25.50	15.0	6.9	*
	_		25.20	15.0	6.9	*
	CV	WR	25.20	15.0	6.2	*
		PR	25.20	13.5	5.8 6.0	*
		ER	25.40	20.0	5.3	*
	E-	CONTROL	25.20	15.0 15.5	6.7	*
	Ea	WR	25.50	13.5	6.4	*
		PR ER	25.50 25.80	20.0	6.2	*
		CONTROL	25.80	16.0	6.6	*
8/19/90		CONTROL	25.00	10.0		
5, 15, 50	Fs	WR	26.00	16.0	6.6	*
		PR	25.50	14.0	6.9	*
		ER	25.50	20.0	6.8	*
		CONTROL	26.00	15.0	6.6	*
	Рp	WR	25.00	15.0	6.6	*
	-		25.00	14.0	5.8	*
		PR	24.70	14.0	5.9	*
			24.70	14.0	6.0	*
		ER	25.20	20.0	6.8	*
			24.70	20.0	6.4	*
		CONTROL	25.00	15.5	6.8	*
			25.00	16.0	6.5	*
	Cv	WR	25.00	15.0	7.7	*
		PR	25.10	14.5	6.5	*
		ER	25.50	20.0	6.7	*
	_	CONTROL	25.00	15.5	6.2	*
	Ea	WR	25.00	15.0	7.6	*
		PR	25.50	14.5	8.2 7.8	*
		ER	25.70	19.5	7.6	*
9/20/00		CONTROL	25.00	15.0	7.0	
8/20/90	Fs	WR	26.00	15.5	3.4	*
	rs	MIX	26.00	15.5	7.5	*
		PR	25.70	14.5	6.8	*
		FK	25.50	14.5	7.7	*
		ER	25.50	20.0	7.7	*
		111	25.50	20.0	7.6	*
		CONTROL	26.00	15.0	6.6	*
			26.00	15.0	6.6	*
	Рp	WR	25.00	15.0	6.7	*
	- F		25.20	15.5	6.4	*
		PR	25.00	15.0	7.0	*
			25.00	15.0	6.5	*
		ER	26.00	20.5	7.2	*

			25.00	20.5	7.6	*
		CONTROL	26.00	15.0	7.0	*
			25.50	15.0	7.2	*
	CV	WR	25.20	15.0	8.5	*
		PR	25.20	15.0	8.3	*
		ER	25.20	20.0	7.9	*
		CONTROL	25.20	15.5	7.0	*
	Ea	WR	25.70	15.0	6.7	*
		PR	25.50	14.5	6.4	*
		ER	26.00	20.0	6.7	*
		CONTROL	25.70	15.5	7.2	*
8/21/90						
- / /	Fs	WR	26.30	15.5	4.9	7.46
		PR	26.00	15.0	7.2	7.80
		ER	26.00	19.0	7.4	7.87
		CONTROL	26.30	15.0	7.3	8.19
	Pp	WR	25.80	14.5	6.4	7.98
	•		25.10	15.5	6.6	7.99
		PR	25.30	14.5	6.2	7.88
		Ti-	25.10	15.0	6.0	7.88
		ER	26.20	20.5	6.8	7.93
			25.50	20.0	6.5	7.84
		CONTROL	26.30	15.0	6.6	8.31
			25.80	15.0	6.3	8.34
	CV	WR	25.20	15.0	8.3	8.27
		PR	25.40	15.0	8.3	8.27
		ER	25.50	18.0	6.7	7.90
		CONTROL	25.30	16.0	6.4	8.25
	Ea	WR	25.50	15.5	6.5	7.94
		PR	25.80	15.0	6.9	8.01
		ER	25.60	19.5	6.3	7.70
		CONTROL	25.80	16.0	7.5	8.31

^{* =} parameter not measured

Table C-2: Experiments conducted with water samples collected from five Potomac River stations from 9/25/90 to 10/2/90. Station abbreviations are as follows: MT= Morgantown, DG= Dahlgren, PP= Possum Point, IH= Indian Head, FP= Freestone Point. Freshwater test species included: Pimephales promelas (Pl), and Ceriodaphnia dubia (Cd). Saltwater test species included: Fundulus spp., Eurytemora affinis, Palaemonetes pugio, and Cyprinodon variegatus. Salinity adjusted test species included: Eurytemora affinis and Palaemonetes pugio.

Date	Test Species	Station	Temp (C)	Sal (ppt)	DO (mg/L)	рН
9/25/9						
3/23/	Fs	MT	25.10	16.5	6.7	7.81
		DG	26.30	19.0	7.2	7.86
		CONTROL	25.50	17.5	6.7	8.04
	Pp	MT	24.20	15.0	6.4	7.82
	-	DG	25.10	19.0	6.1	7.98
		PP	24.10	16.0	6.4	7.97
		IH	23.80	13.0	6.8	7.88
		FP	23.70	15.0	6.4	7.77
		CONTROL	22.70	17.0	7.0	8.00
	Ea	MT	25.20	16.5	6.0	7.75
		DG	25.10	18.0	6.7	7.81
		PP	25.50	15.5	6.8	7.58
		IH	25.30	14.5	6.5 6.5	7.52 7.52
		FP	24.90	14.5	6.2	7.99
		CONTROL	25.40	16.0	6.1	7.08
	Pl	PP	25.20	0 0	6.2	7.19
		IH	25.20 25.10	0	6.0	7.19
		FP		0	6.0	7.53
	cd	CONTROL PP	25.20 25.10	0	8.5	7.08
	Ca	IH	25.10	Ö	8.0	6.38
		FP	25.20	Ö	7.2	6.03
		CONTROL	24.90	Ö	6.9	7.18
9/26/	/an	CONTROL	24.50	· ·	-	
3/20/	Fs	MT	25.90	16.5	7.2	7.98
		DG	25.10	18.0	7.4	7.97
		CONTROL	25.70	16.0	7.4	8.23
	Cv	MT	24.70	21.0	6.8	8.37
	161	DG	24.90	20.0	7.0	8.19
		CONTROL	25.20	20.0	6.3	8.18
	Pр	MT	25.00	17.0	6.3	8.08
	-	DG	25.80	19.0	6.0	8.04
		PP	24.70	16.0	6.2	8.05
		IH	24.40	15.0	6.2	8.02
		FP	24.80	14.5	6.0	8.05
		CONTROL	24.60	17.0	5.8	8.18
	Ea	ТM	24.40	17.5	7.7	7.87

		DG	25.00	19.0	7.0	7.79
		PP	24.50	16.0	6.8	7.73
		IH	24.70	14.5	6.2	7.82
		FP	24.40	14.5	7.2	7.71
		CONTROL	25.20	17.0	7.4	7.86
	Pl	PP	25.10	0	6.2	7.08
		IH	25.20	ŏ	6.3	7.08
		FP	25.20	Ŏ	6.0	7.07
		CONTROL		0	6.1	7.04
	74		25.20			
	Cd	PP	24.90	0	8.2	7.52
		IH	24.80	0	8.0	7.30
		FP	24.80	0	7.9	6.68
		CONTROL	24.40	0	6.7	7.06
9/27/90						
	Fs	MT	26.50	17.0	6.7	7.90
		DG	25.70	18.0	7.2	8.08
		CONTROL	26.70	15.0	7.3	8.32
	CV	MT	25.00	20.0	7.9	8.26
		DG	25.10	18.0	7.8	8.25
		CONTROL	25.10	16.0	7.0	7.97
	Pp	MT	25.00	17.5	6.0	7.97
	- 5	DG	26.20	18.0	6.2	7.95
		PP	25.00	16.0	6.0	7.94
		IH	24.60	15.0	5.8	7.97
					5.3	7.92
		FP	25.00	14.5		
	_	CONTROL	25.00	14.0	7.7	8.21
	Ea	MT	25.00	18.0	6.3	7.98
		DG	25.80	19.0	6.2	7.84
		PP	24.60	16.0	6.0	8.03
		IH	25.10	15.0	5.8	7.88
		FP	24.80	15.0	5.6	7.94
		CONTROL	25.00	15.0	6.6	8.15
	Pl	PP	25.10	0	4.2	7.52
		IH	25.30	0	4.8	7.63
		FP	25.20	0	4.5	7.55
		CONTROL	25.20	Ō	5.6	7.56
	Cd	PP	24.70	Ö	9.3	7.19
	Cu	IH	24.60	Ö	8.2	6.94
					9.0	7.08
		FP	25.00	0	7.9	7.11
0/20/00		CONTROL	25.60	0	7.9	/.11
9/28/90	n -			26.0	4 0	7 70
	Fs	MT	26.40	16.0	4 . 8	7.73
		DG	24.80	18.0	4.9	7.65
		CONTROL	26.10	15.0	4.8	8.06
	CV	MT	25.10	18.0	6.9	8.04
		DG	24.70	19.0	7 • 0	7.98
		CONTROL	25.30	16.0	7 . 1	8.20
	Pр	MT	25.00	16.0	6.5	7.90
	-	DG	25.50	19.0	6.6	7.93
		PP	24.90	16.0	6.9	7.91
		IH	25.30	14.5	7.1	7.91
		FP -		15.0	7.1	7.93
			25.50		7.6	8.17
	E-	CONTROL	25.40	14.6		
	Ea	MT	26.10	16.0	8.3	8.23

	Pl Cd	DG PP IH FP CONTROL PP IH FP CONTROL PP	26.00 26.60 25.90 26.10 25.50 25.00 25.20 25.10 25.10 25.10	17.5 15.5 15.0 15.0 14.5 0 0 0	7.6 8.0 7.5 7.7 7.8 6.7 6.9 7.0 7.0 8.6 8.5	8.16 8.15 8.11 8.18 8.29 7.48 7.37 7.43 7.43 8.21 7.41
		FP	24.40	0	8.4	6.78
9/29/90		CONTROL	25.30	0	8 . 0	7.25
9/29/90	Fs	MT	26.50	15.5	5.4	7.89
		DG	25.40	17.0	5.9	7.80
		CONTROL	26.30	15.0	5.9	8.22
	CV	MT	24.90	16.0	7.1	8.05
		DG	25.30	18.0	7.5	8.09
	Dw	CONTROL	25.00	15.5 16.0	7.5 6.9	8.31 8.00
	Рр	MT DG	25.40 25.10	18.0	6.7	7.99
		PP	24.80	16.0	6.7	8.10
		IH	24.60	15.5	6.5	8.07
		FP	24.70	15.5	7.0	8.14
		CONTROL	24.70	15.0	6.7	8.25
	Ea	MT	25.50	17.0	7.5	8.24
		DG	25.80	16.0	7.7	8.19
		PP	25.00	15.5	7.7	8.18
		IH	25.00	15.0	7.2	8.19 8.38
		FP	25.40	15.0 15.0	7.0 7.5	8.31
	Pl	CONTROL PP	25.60 25.00	0	7.9	7.65
	FI	IH	25.20	Ö	8.4	7.50
		FP	25.20	Ö	7.9	7.58
		CONTROL	25.00	0	8.0	7.95
	Cd	PP	25.30	0	9.4	8.36
		IH	25.80	0	8.2	7.81
		FP	25.80	0	8.4	6.67
		CONTROL	25.10	0	8.2	7.93
9/30/90	D -	1477	26.20	16.0	6.3	7.95
	Fs	MT	26.30	16.0 16.0	6.4	7.93
		DG CONTROL	25.80 26.00	15.0	6.6	8.27
	CV	MT	25.40	16.0	7.2	8.05
	•	DG	25.10	16.0	7 . 2	8.04
		CONTROL	24.60	15.0	7.2	8.33
	Рp	MT	24.50	16.0	7.1	7.99
	-	DĢ	25.50	17.5	7 ∗ 0	7.99
		PP	24.50	16.0	7 1	8.11
		IH	24.60	15.5	7.0	8.08
		FP	24.60	15.5	7.1 7.2	8.06 8.30
	E a	CONTROL	23.60	15.0 16.5	8.0	8.11
	Ea	MT	27.10	16.5	0.0	0.11

		DG PP IH FP	27.30 26.80 27.10 27.20	16.0 15.5 8.0 15.5	7.7 8.0 7.8 7.5	8.00 8.10 7.80 8.10
	Pl	CONTROL PP IH	26.50 25.00 25.70	15.0 0 0	7.6 8.9 8.1	8.31 7.64 7.91
	Cđ	FP CONTROL PP	25.20 24.90 25.10	0 0	8.2 7.1 8.9	7.85 7.76 8.20
	ca	IH FP	25.20 25.70	0 0	9.1 8.7 8.2	8.09 8.09 7.81
10/1/9		CONTROL	25.10	0	0.2	7.01
10/1/9	Fs	MT	25.80 26.10	12.0 12.0	1.7 6.2	7.47 7.93
		DG	25.50	15.5	6.5	7.91
	_	CONTROL	26.10	15.0	6.7 7.5	8.31 8.14
	CV	MT	24.90	17.0 17.0	7.5	8.12
		DG CONTROL	25.20 25.10	15.5	7.5	8.40
	Рp	MT	24.50	17.0	7.5	8.14
	- P	DG	24.60	17.0	7.4	8.12
		PP	24.80	15.5	7.3	8.27
		IH	24.40	15.0	7.3	8.24
		FP	24.30	15.0	7.5	8.29
		CONTROL	24.60	16.0	7.5	8.42
	Ea	MT		DATA	LOST 7.7	8.26
		DG	25.80	17.0 15.0	8.0	8.39
		PP IH	25.80 25.80	11.0	7.8	8.43
		FP	25.90	15.0	8.3	8.35
		CONTROL	26.30	16.0	7.6	8.42
	Pl	PP	25.00	0	8.2	7.95
		IH	25.30	0	8.3	7.96
		FP	25.10	0	7.8	7.91
		CONTROL	25.10	0	7.4	7.45
	Cd	PP	25.10	0	7.0	7.83
		IH	25.50	0	7.8	8.03
		FP	26.20	0	7.1 6.5	7.96 7.81
20/0/0	•	CONTROL	26.20	0	0.5	7.01
10/2/9		мп	26.10	14.0	7.1	7.92
	Fs	MT DG	25.50	15.5	7.1	7.91
		CONTROL	26.10	15.0	7.3	8.23
	CV	MT	24.70	17.0	7.2	7.93
	•	DG	25.20	16.0	6.9	7.90
		CONTROL	24.90	16.0	7.0	8.28
	Pp	MŢ	24.20	17.0	6.9	8.03
		DG	24.60	17.0	7.0	8.04
		PP	24.40	15.0	7.3	8.15 8.16
		IH	24.70	15.0	7.2 7.1	8.08
		FP	24.20	15.0	7.3	8.35
		CONTROL	24.70	16.0	,	0.00

	Ea	MT	25.50	17.0	<10.0	8.68
		DG	25.40	16.0	<10.0	8.56
		PP	22.90	15.0	<10.0	8.77
		IH	24.40	14.0	<10.0	8.83
			24.30	14.0	<10.0	8.93
		FP	25.20	15.0	<10.0	8.75
		CONTROL	25.70	14.0	<10.0	8.68
	Pl	PP	25.50	0	9.7	8.22
		IH	25.70	0	10.2	8.41
		FP	25.30	0	9.6	8.21
		CONTROL	25.30	0	6.8	8.01
	Cd	PP	25.00	0	7.9	8.16
		IH	25.30	0	7.7	8.03
		FP	25.20	0	8.4	8.36
		CONTROL	25.20	0	7.8	8.19
10/3/90						
	CV	MT	25.00	17.0	7.8	8.12
		DG	25.30	16.0	7.4	7.99
		CONTROL	25.50	16.5	7.4	8.32

Table C-3: Laboratory water quality conditions during the second Ceriodaphnia test conducted from 10/27/90 to 11/3/90 using fresh water from the three Potomac River sites.

Date	Station	Temp (C)	DO (mg/L)	рН (Hardness (mg/L CaCO ₃)
10/27	 /90				
20/21/	PP w	23.60	6.5	7.87	92
	IH	23.10	7.0	7.87	92
	FP	22.80	6.9	7.82	84
	CONTROL	22.80	7.2	8.40	136
10/28	/90				
	PP	24.20	7.5	8.15	92
	IH	23.70	7.4	7.95	76
	FP	23.80	7.5	7.98	84
	CONTROL	24.00	7.7	8.48	184
10/29		05 00	7.0	7 07	*
	PP	25.20	7.0	7.97	*
	IH	24.70	7.6	7.93 7.89	*
	FP	24.80	7.6 7.2	8.30	*
10/20	CONTROL	25.60	1.4	0.30	•
10/30	PP	25.30	7.7	7.82	*
	IH	25.10	7.9	7.95	*
	FP	25.50	7.7	7.85	*
	CONTROL	25.10	7.6	8.29	*
10/31					
,	PP	25.10	7.2	7.95	*
	IH	25.00	7.2	8.15	*
	FP	25.20	7.3	8.07	*
	CONTROL	24.70	7.5	8.36	*
11/1/	′90				
	PP	26.60	7.4	7.97	*
	IH	26.50	7.7	7.90	*
	FP	26.00	7.9	7.87	*
	CONTROL	26.00	7.6	8.30	*
11/2/	/90				_
	PP	24.50	7 , 3	7.90	*
	IH	25.20	7.2	7.88	*
	FP	25.00	7.7	7.91	*
	CONTROL	24.60	7.7	8.26	*
11/3/			<u> </u>	7 07	*
	PP	24.80	7 4	7.97	*
	IH	24.80	7.6	7.98	*
	FP	24.80	7.4	7.97	*
	CONTROL	24.80	7.5	8.40	~

^{* =} parameter not measured

Appendix D

Organic analysis data are presented for
Possum Point, Freestone Point, Dahlgren,
Elizabeth River and the Nansemond River sediments.
The semi-volatile organic compounds were identified and quantitated against a user-created "priority pollutants" library which matches both retention times and spectral fit.
The quantitations were based upon 5-7 point calibration curves.
The tentatively identified compounds were identified by spectral fit against the NBS library and quantitated against an internal standard, assuming a 1:1 response factor.

w² 2° \$

Table D-1 Organics analysis data sheet for semi-volatile compounds.

Laboratory: Organics Contractor: MD DNR Project ID: Ambient Toxicity Sample #: 35363

Sample ID: Possum Point

Dates: Received - 9/26/90 Extracted - 11/21/90 Analyzed - 11/30/90

Method: EPA 3550/8270 Instrument: INCOS50

Analyst: TLP Data Released By: M. Helmstetter

Matrix: Sediment Units: μ g/kg dry Sample w/v: 30.04 % Moisture: 49.52

CAS #	Compound	Concentration	Detection Limit
84-74-2	Di-n-butylphthalate	77 *	5.9
129-00-0	Pyrene	65	10.6
218-01-9	Chrysene	186	14.5
117-81-7	Bis(2-ethylhexyl)phthalate	670	12.5
50-32-8	Benzo(a)pyrene	155	15.2

^{*} Compound detected in QC blank:

Organics analysis data sheet for tentatively identified semi-volatile compounds. Table D-2

Contractor: MD DNR Laboratory: Organics Project ID: Ambient Toxicity

Sample #: 35363 Sample ID: Possum Point

Extracted - 11/21/90 Received - 9/26/90 Dates:

Analyzed - 11/30/90

Instrument: INCOS50 Method: EPA 3550/8270

Data Released By: M. Helmstetter Analyst: TLP

Units: $\mu g/kg dry$ Matrix: Sediment % Moisture: 49.52 Sample w/v: 30.04

21.50.0	<u> </u>	Scan #	Estimated Concentration*
CAS #	Compound	Scall #	Civil Control Civil
	Unknown	511	4310 **
75-91-2	Hydroperoxide, 1,1-dimethylethyl	516	675 **
75-71-2	Unknown	529	635 **
4160-75-2	2-propanone, 1-cyclopropyl-	535	369 **
75-91-2	Hydroperoxide, 1,1-dimethylethyl	582	11200 **
15-71-2	Unknown	592	16300 **
	Unknown	635	223000
4305-26-4	2-hexanone, 6-(acetyloxy)-	756	10800
17257-81-7	Ethanone, 1-(3-ethyloixeahyl)-	786	937
3240-09-3	5-hexen-2-one, 5-methyl	843	5780
3240 07 3	Unknown	867	1830
26118-38-7	2-hexanone, 3,3-dimethyl-	931	2900
56052-85-8	2-pentene, 5-(pentyloxy)-, (E)-	954	12800
5343-96-4	2-butanol, 3-methyl-, acetate	999	3750
542-59-6	1,2-ethanediol, monoacetate	1114	6360
10544-50-0	Sulfur, mol (S8)	2896	6690
3891-98-3	Dodecane, 2,6,10-trimethyl-	4139	1400
3891-98-3	Dodecane, 2,6,10-trimethyl-	3938	842
6971-40-0	17-pentatriacontene	3949	871
107-41-5	2.4-pentanediol, 2-methyl	1420	780
107-41-5	2.4-pentanediol, 2-methyl	1420	/60

^{*} Estimated concentration is based on a 1:1 response with the internal standard.

^{**} Compound detected in QC blank.

Table D-3 Organics analysis data sheet for semi-volatile compounds.

Laboratory: Organics Contractor: MD DNR
Project ID: Ambient Toxicity Sample #: 35362

Dates: Received - 9/26/90
Analyzed - 11/30/90

Contractor: MD DNR
Sample #: 35362

Extracted - 11/21/90

Method: EPA 3550/8270 Instrument: INCOS50

Analyst: TLP Data Released By: M. Helmstetter

Matrix: Sediment Units: μ g/kg dry Sample w/v: 30.08 Units: μ g/kg dry % Moisture: 49.09

CAS #	Compound	Concentration	Detection Limit
218-01-9	Chrysene Bis(2-ethylhexyl)phthalate	210	14.5
117-81-7		490 *	12.5

^{*} Compound detected in QC blank.

Table D-4 Organics analysis data sheet for tentatively identified semi-volatile compounds.

Laboratory: Organics Contractor: MD DNR
Project ID: Ambient Toxicity Sample #: 35362

Sample ID: Freestone Point

Dates: Received - 9/26/90 Extracted - 11/21/90

Analyzed - 11/30/90

Method: EPA 3550/8270 Instrument: INCOS50

Analyst: TLP Data Released By: M. Helmstetter

Matrix: Sediment Units: μ g/kg dry Sample w/v: 30.08 Units: μ g/kg dry % Moisture: 49.09

		Estimated
Compound	Scan #	Concentration*
** 1		
		3060
2-propyn-1-01, acetate	505	1120 **
Unknown	543	1020 **
Hydroperoxide, 1,1-dimethylethyl	563	22000
2-propyn-1-01, acetate	613	162000 **
2-hexanone, 6-(acetyloxy)-	744	7460
Hexane, 2-bromo-	777	723
	836	4880
•	860	1820
Unknown	924	2100
2-pentene, 5-(pentyloxy)-, (E)-	949	13200
	962	8000
•	994	3500
	1116	4890
Unknown	1005	621
Sulfur, mol. (S8)	2890	4020
	3938	804
		657
		1030
		647
	Unknown 2-propyn-1-01, acetate Unknown Hydroperoxide, 1,1-dimethylethyl 2-propyn-1-01, acetate 2-hexanone, 6-(acetyloxy)- Hexane, 2-bromo- 5-hexen-2-one, 5-methyl- 2-propyn-1-01, acetate Unknown 2-pentene, 5-(pentyloxy)-, (E)- 3-heptanone, 2,4-dimethyl 3-heptanone, 2,4-dimethyl 2,3-butanedione	Unknown 451 2-propyn-1-01, acetate 505 Unknown 543 Hydroperoxide, 1,1-dimethylethyl 563 2-propyn-1-01, acetate 613 2-hexanone, 6-(acetyloxy)- 744 Hexane, 2-bromo- 777 5-hexen-2-one, 5-methyl- 836 2-propyn-1-01, acetate 860 Unknown 924 2-pentene, 5-(pentyloxy)-, (E)- 949 3-heptanone, 2,4-dimethyl 962 3-heptanone, 2,4-dimethyl 994 2,3-butanedione 1116 Unknown 1005 Sulfur, mol. (S8) 2890 Undecane, 3,8-dimethyl 3938 17-pentatriacontene 3951 Decanedioic acid, didecyl ether 4138

^{*} Estimated concentration is based on a 1:1 response with the internal standard.

^{**} Compound detected in QC blank.

Table D-5 Organics analysis data sheet for semi-volatile compounds.

Laboratory: Organics Contractor: MD DNR
Project ID: Ambient Toxicity Sample #: 35360
Sample ID: Dahlgren

Dates: Received - 9/26/90 Extracted - 11/21/90 Analyzed - 11/30/90

Method: EPA 3550/8270 Instrument: INCOS50

Analyst: TLP Data Released By: M. Helmstetter

Matrix: Sediment Units: μ g/kg dry Sample w/v: 30.00 % Moisture: 27.06

CAS #	Compound	Concentration	Detection Limit
56-55-3	Benzo(a)anthracene Bis(2-ethylhexyl)phthalate	99.8	17.8
117-81-7		691 *	12.5

^{*} Compound detected in QC blank.

Table , D-6 Organics analysis data sheet for tentatively identified semi-volatile compounds.

Laboratory: Organics

Ambient Toxicity

Contractor: MD DNR

Project ID: Sample ID:

Dahlgren

Sample #: 35360

Dates:

Received - 9/26/90

Analyzed - 11/30/90

Extracted - 11/21/90

Method: EPA 3550/8270

Analyst: TLP

Instrument: INCOS50

Data Released By: M. Helmstetter

Matrix: Sediment Sample w/v: 30.00

Units: $\mu g/kg dry$ % Moisture: 27.06

			Estimated
CAS #	Compound	Scan #	Concentration*
563-80-4	2-butanone, 3-methyl-	790	2230
20019-64-1	2(5H)-furanone, 5,5-dimethyl	839	620
	Unknown	865	341
	Unknown	930	1660
18641-71-9	3-heptanone, 2,4-dimethyl-	966	5660
18641-71-9	3-heptanone, 2,4-dimethyl-	1001	8060
	Unknown	1012	553
	Unknown	1094	955
431-03-8	2,3-butanedione	1111	230
10544-50-0	Sulfur, mol. (S8)	2886	1270
74630-61-8	2-undecene, 6-methyl-, (E)-	3106	1270

Estimated concentration is based on a 1:1 response with the internal standard.

Table D-7 Organics analysis data sheet for semi-volatile compounds. (Note: Double underlined values represent concentrations exceeding "Effects Range-Medium" levels for selected polynuclear aromatic hydrocarbons, as defined in Long and Morgan, 1990).

Laboratory: Organics
Project ID: Ambient Toxicity
Sample ID: Atlantic Wood

Dates: Received - 8/16/90

Contractor: MD DNR
Sample #: 34927

Extracted - 12/14/90

Analyzed - 12/14/90

Method: EPA 3550/8270 Instrument: INCOS50

Analyst: TLP Data Released By: M. Helmstetter

Matrix: Sediment Units: μg/kg dry Sample w/v: 30.00 % Moisture: 62.7

Detection Limit Concentration Compound CAS # 4.6 43.7 91-20-3 Naphthalene 5.9 486 208-96-8 Acenaphthalene 3.11x10 ³ 9.9 Acenaphthene 83-32-9 7.9 486 Dibenzofuran 132-64-5 9.9 <u>690</u> Fluorene 86-73-7 <u>5.31x10</u> ³ 9.2 85-01-8 Phenanthrene 7.10×10^{-3} 9.9 120-12-7 Anthracene 1.07x10 4 10.6 Fluoranthene 206-44-0 1.18x10 ⁴ 10.6 Pyrene 129-00-0 9.18x10 3 14.5 Chrysene 218-01-9 12.5 146 * Bis(2-ethylhexyl)phthalate 117-81-7 $6.29x10^{-3}$ 13.9 Benzo(b)fluoranthene 205-99-2 6.23x10³ 15.2 Benzo(a)pyrene 50-32-8 2.16x10 17.8 Dibenzo(a,h)anthracene 53-70-3 3.42x10 ³ 16.5 Indeno(1,2,3-cd)pyrene 193-39-5 3.89x10³ 16.5 191-24-2 Benzo(g,h,i)perylene

^{*} Compound detected in QC blank.

Table D-8 Organics analysis data sheet for tentatively identified semi-volatile compounds.

Laboratory: Organics Contractor: MD DNR
Project ID: Ambient Toxicity Sample #: 34927

Sample ID: Atlantic Wood

Dates: Received - 8/16/90 Extracted - 12/14/90

Analyzed - 12/14/90

Method: EPA 3550/8270 Instrument: INCOS50

Analyst: TLP Data Released By: M. Helmstetter

Matrix: Sediment Units: μ g/kg dry Sample w/v: 30.00 % Moisture: 62.7

			Estimated
CAS #	Compound	Scan #	Concentration*
613-12-7	Anthracene, 2-methyl-	2690	4750
	Unknown	2714	2960
203-64-5	4H-cyclopenta(def)phenanthrene	2722	11200
13141-45-2	Benzene, 1,1'-(1-buten-3-yne-1,4-diyl)bis-	2803	3130
129-00-0	Pyrene **	2942	24800
129-00-0	Pyrene **	3013	21900
243-42-5	Benzo(b)naphthol[2,3-D]furan	3020	5730
238-84-6	11H-benzo(a)fluorene	3143	12300
2381-21-7	Pyrene, 1-methyl	3166	9200
205-43-6	Benzo(b)naphthol[1,2-D]thiophene	3354	2890
195-19-7	Benzo(c)phenanthrene	3368	4890
217-59-4	Triphenylene	3464	11900
	Unknown	3482	4910
2871-91-2	Triphenylene, 1-methyl	3595	2980
20/1/12	Unknown	3639	4590
	Unknown	3842	2840
886-38-4	2-cyclopropen-1-one, 2,3-diphenyl	3880	4390
205-82-3	Benzo(j)fluoranthene	3911	10700
192-97-2	Benzo(e)pyrene	3936	2800
174-71-4	Unknown	3987	3850
	Chritowii	5701	

^{*} Estimated concentration is based on a 1:1 response with the internal standard.

^{**} Compound name based on NBS library match. These compounds do not have the correct retention time for Pyrene which is found in this sample at its correct retention time.

Table D-9 Organics analysis data sheet for semi-volatile compounds.

Laboratory: Organics Contractor: MD DNR
Project ID: Ambient Toxicity Sample #: 35378

Dates: Collected - 9/25/90 Received - 9/27/90
Extracted - 11/21/90 Received - 11/30/90

Method: EPA 3550/8270

Analyst: TLP

Matrix: Sediment Sample w/v: 30.01 Instrument: INCOS50
Data Released By: M.

Data Released By: M. Helmstetter

Units: μg/kg dry % Moisture: 51.95

CAS #	Compound	Concentration	Detection Limit
117-81-7	Bis(2-ethylhexyl)phthalate Benzo(a)anthracene	32 *	12.5
56-55-3		193	17.8

^{*} Compound detected in QC blank.

Table D-10 Organics analysis data sheet for tentatively identified semi-volatile compounds.

Laboratory: Organics Contractor: MD DNR
Project ID: Ambient Toxicity Sample #: 35378

Sample ID: Nansemond Reference

Dates: Collected - 9/25/90 Received - 9/27/90 Extracted - 11/21/90 Analyzed - 11/30/90

Method: EPA 3550/8270 Instrument: INCOS50

Analyst: TLP Data Released By: M. Helmstetter

Matrix: Sediment Units: μ g/kg dry Sample w/v: 30.01 % Moisture: 51.95

			Estimated
CAS #	Compound	Scan #	Concentration*
	Unknown	451	2200
75-91-2	Hydroperoxide, 1,1-dimethylethyl	516	15600
	Unknown	555	848
75-91-2	Hydroperoxide, 1,1-dimethylethyl	585	24600
	Unknown	629	298000
	Unknown	752	8910
	Unknown	784	1700
758-87-2	4-penten-2-one, 3-methyl	838	3050
	Unknown	862	1680
18641-71-9	3-heptanone, 2,4-dimethyl	962	7420
18641-71-9	3-heptanone, 2,4-dimethyl	996	5600
	Unknown	1007	762
542-59-6	1,2-ethenediol, monoacetate	1102	1290
	Unknown	1417	425
10544-50-0	Sulfur, mol. (S8)	2881	920

^{*} Estimated concentration is based on a 1:1 response with the internal standard.

Appendix E

Pesticide analysis data for Possum Point, Freestone Point, Dahlgren, Elizabeth River and Nansemond River sediments

with

Table E-1 Organics analysis data sheet for tentatively identified pesticide compounds.

Laboratory: Organics
Project ID: Ambient Toxicity

Sample ID: Possum Point

Dates: Received - 9/26/90

Analyzed - 1/14/91

Method: EPA 3550/8080

Analyst: RJM

Matrix: Sediment Sample w/v: 30.02 Instrument: GC-6

Extracted - 12/14/90

Contractor: MD DNR

Sample #: 35363

Data Released By: M. Helmstetter

Units: μ g/kg dry % Moisture: 49.52

240 "	Company	Concentration	Detection Limit
CAS #	Compound	Concentration	
72-55-9	4,4'-DDE	1.05	0.528

Table E-2 Organics analysis data sheet for tentatively identified pesticide compounds.

Contractor: MD DNR Laboratory: **Organics** Project ID: Ambient Toxicity Sample #: 35362 Sample ID: Freestone Point Received - 9/26/90 Extracted - 12/14/90 Dates: Analyzed - 1/14/91 Method: EPA 3550/8080 Instrument: GC-6 Data Released By: M. Helmstetter Analyst: RJM

Matrix: Sediment Units: μ g/kg dry Sample w/v: 30.06 % Moisture: 49.09

CAS #	Compound	Concentration	Detection Limit
391-84-6	α-BHC	0.574 *	0.714
72-55-9	4,4'-DDE	0.353 *	0.528

^{*} Compound detected below calculated method limit.

Table E-3 Organics analysis data sheet for tentatively identified pesticide compounds.

Laboratory: Organics Contractor: MD DNR
Project ID: Ambient Toxicity Sample ID: Dahlgren

Dates: Received - 9/26/90
Analyzed - 1/14/91

Contractor: MD DNR
Sample #: 35360

Extracted - 12/14/90

Method: EPA 3550/8080

Analyst: RJM

Matrix: Sediment Sample w/v: 30.03 Instrument: GC-6

Data Released By: M. Helmstetter

Units: μ g/kg dry % Moisture: 27.06

CAS #	Compound	Concentration	Detection Limit
1024-57-3	Heptachlor epoxide	2.06	0.570

Table E-4 Organics analysis data sheet for tentatively identified pesticide compounds. (Note: Single underlined values represent concentrations exceeding the "Effects-Range Low"; double underlined values represent concentrations exceeding "Effects Range-Medium" levels for selected chlorinated pesticides, as defined in Long and Morgan, 1990.)

Laboratory: Organics Contractor: MD DNR
Project ID: Ambient Toxicity Sample #: 34927
Sample ID: Atlantic Wood

Dates: Received - 8/16/90 Extracted - 12/14/90 Analyzed - 1/14/91

Method: EPA 3550/8080 Instrument: GC-6

Analyst: RJM Data Released By: M. Helmstetter

Matrix: Sediment Units: μ g/kg dry Sample w/v: 30.06 % Moisture: 62.7

CAS #	Compound	Concentration	Detection Limit
391-84-6	α-ВНС	3.67	0.714
391-86-8	δ-BHC	6.59	1.05
1024-57-3	Heptachlor epoxide	5.57	0.570
959-98-8	Endosulfan I	1.88	0.859
72-55-9	4,4'-DDE	11.8	0.528
60-57-1	Dieldrin	<u>29.7</u>	0.898
7421-93-4	Endrin aldehyde	2.16	2.41

Table E-5 Organics analysis data sheet for tentatively identified pesticide compounds.

Laboratory: Organics

Project ID: Ambient Toxicity

Sample ID: Nansemond Reference

Dates:

Received - 9/25/90

Analyzed - 1/14/91

Method: EPA 3550/8080

Analyst: RJM

Matrix: Sediment Sample w/v: 30.09 Contractor: MD DNR

Sample #: 35378

Extracted - 12/14/90

Instrument: GC-6

Data Released By: M. Helmstetter

Units: μg/kg dry % Moisture: 51.95

CAS # Compound Concentration Detection Limit

No compounds detected

					2
					71
		3			
					2
		38			
	(d)		2		
				8	
	(*) (K		OF 11	8. 2	