DRAFT FINAL MPRSA Section 103 Sediment Characterization Testing and Analysis San Juan Harbor, Puerto Rico Contract: W912PM-15-D-0006 Order Number: W912EP-21-F-0026 Submitted to U.S. Army Corps of Engineers Jacksonvi District 701 San N oulevard Jackson ville, Florida 32207-8175 Prepared by: AMAR Environmental Consulting, Inc. 2106 NW 67th Place, Suite 5 Gainesville, FL 32653 www.anamarinc.com Environmental Consulting, Inc. April 2021



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APPENDICES

NOTE: Appendices (with the exception of Appendix G) are provided in electronic format only and may be found on the accompanying disc.

- Appendix A SAP/QAPP
- Appendix B Field Paperwork
- Appendix C Sediment Physical Lab Report
- Appendix D Chemistry Lab Reports (Sediment, Elutriate/Site Water, Tissues)
- Appendix E Chemical Quality Assurance Report
- Appendix F Statistical Calculations
- Appendix G Toxicology Lab Report
- Appendix H ADDAMS Model Files
- Appendix I Photos of Samples
- Appendix J Pertinent Correspondence



ACRONYMS, ABBREVIATIONS, AND INITIALISMS

	internet and the internet inte
ADDAMS	Automated Dredging and Disposal Alternatives Modeling System
ARI	Analytical Resources, Inc.
CCV	continuing calibration verification
CETIS	Comprehensive Environmental Toxicity Information System
CFR	Code of Federal Regulations
CMC	criteria maximum concentration (synonymous with 'acute')
CQAR	Chemical Quality Assurance Report
DQCR	Daily Quality Control Report
EC ₅₀	effective concentration affecting 50% of a population
ECD	electron capture detector
EPA/USEPA	U.S. Environmental Protection Agency
ERL	effects range-low
FDA	U.S. Food and Drug Administration
GC/MS	gas chromatography/mass spectrometry
GC-ECD	gas-chromatography-electron capture detection
HMW	high molecular weight
ICP/MS	inductively coupled plasma/mass spectrometry
ICV	initial calibration verification
ITM	Inland Testing Manual (EPA and USACE 1998)
LC ₅₀	lethal concentration 50%
LCS	laboratory control sample
LMW	low molecular weight
LPC	limiting permissible concentration
MDL	method detection limit
mg/L	milligrams per liter
MLLW	mean lower low water
MLW	mean low water
MPRSA	Marine Protection, Restarshand Sinctuaries Act
MRL	method reporting limit
MRM	multiple reaction monitoring
NOAA	National Oceanis and Atmospheric Administration
NOAA	no-observed affect oncentration
NU	
	nephelometric tradiction it
ODMDS	ocean dredged material disposal site
PAH	polycycli (arc n tic hydrocarbon
PCB	polychonnated aphenyl
ppt	parts for the band
QA	que lity ansurance
QAPP	Quary Assurance Project Plan
QC	quality control
RPD	telanve percent difference
RTM	(MY Army Corps of Engineers District) Regional Testing Manual
SAP/OAP 2	Sampling and Analysis Plan/Quality Assurance Project Plan
SD 5	sodium dodecyl sulfate
SOP P	standard operating procedure
RRM	standard reference material
	threshold effects level
TOC	total organic carbon
SACE	U.S. Army Corps of Engineers
	U.S. Soil Classification Systems
▼	



EXECUTIVE SUMMARY

This report details the field sampling, analysis, and results of MPRSA Section 103 sediment testing and analysis in support of the San Juan Harbor dredging operations. Sampling and testing were performed for both maintenance and new work dredging to allow for deepening and widening of the channels within the San Juan Harbor. Field sampling, compositing, and shipping preparations took place on October 19 through November 2, 2020.

Areas proposed to be dredged have been divided into reaches or dredging units. The rational for the sampling approach is summarized in Section 2.1. All samples within each dredging unit were collected either by vibracore to project depth or refusal or by a grab sampler. Samples within each dredging unit were composited and homogenized to create one composite per dividging unit: Analysis of the composited sediment consisted of three analytical tiers: physical created, and toxicological/bioaccumulation.

Sediment Physical Results

Grain size distribution and total solids were analyzed in project composite samples, subsamples, individual clay/native material samples, and the reference sample. The following parameters were also analyzed for the composite sample: bulk density, specific gravity, and Atterberg limits. Grain size results for the composite and clay/native material samples are summarized below.

San Juan Harbor Maintenance Reach A

M-A-S-20-COMP was composed primarily of silt/clay (86.4%) with 13.6% sand.

San Juan Harbor Maintenance Reach B

M-B-S-20-COMP was composed primerily of silt/cay (68.1%) with 31.9% sand.

Army Terminal Widener Reach

D-ATw-20-COMP was composed primarily assilt/clay (78.2%) with 20.8% sand and 1.0% gravel.

San Antonio Extension

D-SAx-20-COMP was composed rimarily of silt/clay (85.3%) with 14.1% sand and 0.6% gravel.

Individual Clay/Native Material Samples

Sample D-EC-C 2 (colocated with station M-A-S-3) from the Entrance Channel in Reach A was primarily corposed of fine material with 96.7% silt/clay with 3.3% sand.

Sample D-AN-C-1 co-located with station D-ATw-S-1) in the Army Terminal Widener was primarily compound of fine material with 83.4% silt/clay with 15.2% sand and 1.4% gravel.

Sample L ATw-C-2 (co-located with station D-ATw-S-2) in the Army Terminal Widener was primary composed of fine material with 66.0% silt/clay with 34% sand.

ample D-ATw-C-3 (co-located with station D-ATw-S-4) in the Army Terminal Widener was repainly composed of fine material with 54.7% silt/clay with 45% sand.

Sample D-ATw-C-4 (co-located with station D-ATw-S-3) in the Army Terminal Widener was primarily composed of fine material with 59.5% silt/clay with 39.1% sand and 1.4% gravel.

Reference

SJH20-REF was primarily composed of silt and clay (90.1%) with 9.1% sand.



Sediment Chemistry

Sediment composites, clay/native material samples, and the reference (SJH20-REF) were analyzed for total solids, TOC, metals, pesticides, PAHs, and PCBs. The subsamples were also analyzed for total solids and TOC. Comparisons of sediment chemistry results were made to the TEL and ERL, where available.

<u>Metals</u>

All nine metals analyzed were detected in concentrations above the MDL in all of the aroust composite samples. With the exception of cadmium, all other metals analyzed were also detecte in concentrations above the MDL in the reference and several of the individual clay/native metals subsamples. Concentrations of metals were below applicable TEL and ERL thresholds with the exceptions summarized below.

Composite Samples

- M-A-S-20-COMP: arsenic, copper, mercury and nickel exceeded her FL and (or) ERL.
- M-B-S-20-COMP: arsenic, copper, and mercury exceeded the TEL and (or) ERL.
- D-ATw-S-20-COMP: arsenic, copper, and nickel exceeded the TM, and (or) ERL.
- D-SAx-S-20-COMP: arsenic, copper, lead, mercury, nick 1 silver, and zinc exceeded the TEL and (or) ERL.

Clay/Native Material Samples

- D-EC-C-2: arsenic, copper, mercury, and nickel acceeded the TEL and (or) ERL.
- D-ATw-C-1, C-2, and C-4: arsenic and corper exceeded the TEL and (or) ERL.
- D-ATw-C-3: copper exceeded the TEL.

Reference

SJH20-REF had concentrations of arso ic, somer, and nickel that exceeded the TEL and (or) the ERL.

Pesticides

Two of the 15 pesticides tested (p,p')(2,4')-DDE and p,p'(4,4')-DDE] were detected above the MDL (J-qualified or great r) is one or more samples. Concentrations of pesticides were below applicable TEL and EPL meshods with the exceptions summarized below.

Composite Sumpl

• D-Sxx-S-20-COMP: p,p' (4,4')-DDE concentrations exceeded the ERL and TEL.

Clay/Native Material Samples

None of the pesticides were detected in concentrations greater than the MDL for any of the clay same same same u-qualified.

Befe en e

Note of the results for SJH20-REF were detected in concentrations greater than the MDL; all results were U-qualified.

PAHs

All 16 PAH analytes tested were detected above the MDL (J-qualified or greater) in one or more composites or subsamples. Concentrations of PAHs were below applicable TEL and ERL thresholds with the exceptions summarized below.



Composite Samples

- M-A-S-20-COMP, M-B-S-20-COMP, and D-SAx-20-COMP: acenaphthylene and dibenzo(a,h)anthracene concentrations exceeded the TEL.
- D-SAx-20-COMP, benzo(a)pyrene and total HMW PAHs concentrations exceeded the TEL.

Clay/Native Material Samples

None of the results exceeded the TEL or ERL.

Reference

None of the results for SJH20-REF exceeded the TEL or ERL.

<u>PCBs</u>

Up to 20 of the 22 PCB congeners tested were detected in concentration (ab) we the MDL in one or more samples. Concentrations of PCBs were below applicable TEL an EFL thresholds with the exceptions summarized below.

Composite Samples

All composite samples had total EPA Region 2 PCB concentrations that exceeded the TEL and ERL.

Clay/Native Material Samples

All clay/native material samples had total EPA Region 2 PCB concentrations that exceeded the TEL and (or) ERL.

Reference

None of the 22 PCB congeners were between in concentrations greater than the MDL (Uqualified) in SJH20-REF. The reference had total EPA Region 2 PCB concentrations that exceeded the TEL.

Elutriate and Water Chemi

Site water (SJH20-SW), (effects ce water (SJH20-REF-SW), and elutriates generated from the four project composite over saralyzed for metals, pesticides, and PCBs. Results for elutriate and water samples are compared to the CMC from EPA (2006, 2015).

Metals

None of the metals analyzed were detected in concentrations greater than the CMC in any elutriate or water sample.

Per de es

Note of the pesticides analyzed were detected in concentrations greater than the CMC or MDL many platriate or site water samples (U-qualified).

PCBs

None of the PCB congeners were detected in concentrations greater than the MDL in any elutriate or site water samples (U-qualified). There are no CMCs for the PCB congeners tested.



Toxicology

Benthic Bioassays

Significant benthic toxicity, relative to the reference treatment, was observed in the *A. abdita* amphipod test for test sample D-ATw-S-20-COMP only. No significant toxicity was observed in *A. bahia* mysid test. Mean percent survival in the project composite samples was within the specific test criterion (20% of the reference: amphipod; 10% of the reference: mysid), indicating that the test treatments met the LPC for disposal for these tests.

Water Column Bioassay

No statistically significant toxicity was observed in the 100% elutriate concentrations or the A. bahia, M. beryllina, and M. galloprovincialis tests.

Bioaccumulation Potential

No significant toxicity was observed in the bioaccumulation tests. Survival in the reference and test treatment was ≥93.0%, suggesting that adequate tissue mass was available for chemical analyses.

Tissue Chemistry

Wet weight tissue chemistry results for the four project samples are oppared to the reference (SJH20-REF) and to applicable FDA action levels from FDI (2001, 2011).

Lipids and Total Solids

M. nasuta – Total solids ranged from 16.34% to 18.62% along the project samples, reference, and pre-exposure tissues. Lipids ranged from 1.5% to 2.5% among these samples.

A. virens – Total solids ranged from 14.06% to 11.68% among the project samples, reference, and pre-exposure tissues. Lipids ranges from 2.1% to 3.6% among these samples.

<u>Metals</u>

M. nasuta – All metals tested usive ontected in concentrations greater than the MRL in the project samples and the reference. Mean concentrations of lead in the project sample M-B-S-20-COMP were statistically significantly reater than those of the reference. Mean concentrations of lead, silver, and zinc in the project sample D-SAx-S-20-COMP were statistically significantly greater than those of the reference of the reference applicable project sample D-SAx-S-20-COMP were statistically significantly greater than those of the reference of the reference applicable project.

A. virens – Although ested were detected in concentrations greater than the MRL in the project samples and the operance. Mean concentrations of arsenic, cadmium, and chromium in all four project samples were statistically significantly greater than those of the reference. In addition, mean concentrations of copper, nickel, and zinc were statistically significantly greater in D-ATw-S-20-20-4P than those of the reference. None of the mean concentrations of metals exceeded polliable FDA action levels.

Pesticides

M. nasuta – With the exception of 4,4'-DDE in sample D-SAx-S-20-COMP, none of the pesticides were detected in concentrations greater than the MDL in any of the project samples or reference (U-qualified). Mean concentration of 4,4'-DDE (1.49 μ g/kg) in sample D-SAx-S-20-COMP was statistically significantly greater than that of the reference (0.14 μ g/kg). None of the mean concentrations of pesticides exceeded applicable FDA action levels.



A. virens – None of the pesticides were detected in concentrations greater than the MDL in the project samples or the reference. All results were U-qualified. The MDL and MRL for transnonachlor were elevated above the target detection limit due to matrix interference. None of the mean concentrations of pesticides exceeded applicable FDA action levels.

<u>PAHs</u>

M. nasuta – None of the PAHs were detected in concentrations greater than the MDL in the project samples or the reference. All results were U-qualified; therefore, no further statistical analyses comparisons were needed.

A. virens – None of the PAHs were detected in concentrations greater than the MDL in the project samples or the reference. All results were U-qualified; therefore, no further statistical values or comparisons were needed.

Polychlorinated Biphenyls (PCBs)

M. nasuta – Nine of the PCB congeners tested were detected above the MRL in at least one of the project sample replicates. Concentrations of PCB congeners 49, 52, 101, 118, 138, and 153 and total EPA Region 2 PCBs in some of the project samples were statistically significantly greater than those of the reference, as summarized below. Total EFA Popion 2 PCB mean concentration in the project samples did not exceed the FDA action level.

- M-A-S-20-COMP PCB 153
- M-B-S-20-COMP PCBs 49, 52, 153, and Total PCBs
- D-SAx-S-20-COMP PCBs 49, 52, 101, 18, 138, 153 and Total PCBs

A. virens – Nine of the PCB congeners tested were detected above the MRL in at least one of the project sample replicates. Concentrations on ICP congeners 49, 52, 101, and total EPA Region 2 PCBs in some of the project samples were statistically significantly greater than those of the reference, as summarized below. Total EPA Region 2 PCB mean concentration in the project samples did not exceed the FDA action level.

- M-A-S-20-COMP PCB
- M-B-S-20-COMP POR 10
- D-SAx-S-20-COMP PDBs 49, 52, 101, and Total PCBs

ADDAMS V odel

STFATE modeling was performed using two types of dredging equipment, a clamshell dredge combined with a secarate barge or scow and a hopper or cutter dredge. Each type of dredging equipment was modeled with a capacity of 4,800 cubic yards per load based on the largest option currently available in Puerto Rico. The model was also performed with a volume of 15,000 cubic yards per load based on the largest option currently available in case a larger dredging vessel becomes available. All model runs met the disporal ordering for both dredging methods and volumes. Therefore, the material may be on second without location or volume restrictions, to a maximum volume of 15,000 cubic yards per load within the ODMDS boundaries in accordance with all criteria specified by EPA Region 2 and ISACE Jacksonville District.



1 INTRODUCTION

1.1 Project Area Description

The sediment characterization and testing performed for this project includes both routine maintenance material from the San Juan Harbor navigation channels to authorized depths and proposed deepening (new work) material in support of future deepening and widening in some areas of San Juan Harbor. This report summarizes the results of the sampling and testing performed to determine the suitability of the material for disposal in the San Juan Harbor ocenn dredged material disposal site (ODMDS).

Exhibit 1-1 provides an overview of the planned improvements to the San Juan Habor Federal Navigation Project. Harbor improvements, as described in the project work score are broadly described below.

- 1) Deepening of Bar and Entrance Channels to various depths ranging tom -56 to -44 feet,
- 2) Deepening of the Anegado and Army Terminal Channels to -44 fet + (-2 feet overdepth)
 = -46 feet
- 3) Deepening of the San Antonio Channel and San Antonio Application Channel to -36 feet + (- 2 feet overdepth) = -38 feet
- 4) Widening of Army Terminal Channel, and
- 5) Extending the San Antonio Channel in an easter direction.

The project area was divided into five dredging units of reaches for sampling and testing purposes: two maintenance reaches (Reach A and Reach B) and three deepening/widening reaches (Army Terminal Widener; San Antonia Extension; Deepening Reach). In addition, individual samples of clay or native materia were sampled at various locations. These data will be used to compare results from testing conducted in 2000. A detailed description of each reach and information on the sampling and compasiting plan are provided in Section 2. Exhibit 1-2 provides an overview of the target sampling locations for each dredging unit/reach.

MPRSA Section 103 Sediment Characterization San Juan Harbor, Puerto Rico



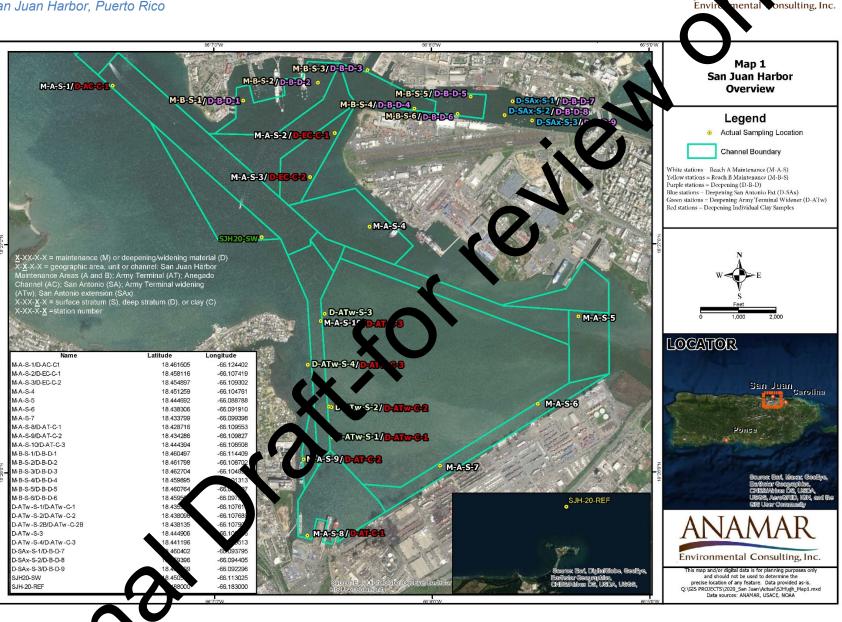


Ex (ibi) 1.

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Planned Improvements to the San Juan Harbor Federal Navigation Project (from Project Work Scope 2020)

MPRSA Section 103 Sediment Characterization San Juan Harbor, Puerto Rico



hibit 1-2. Overview of Project Dredging Units/Reaches and Sampling Locations



1.2 Description of the Testing Approach

1.2.1 Evaluation of Dredge Materials for Disposal

Sediment and suspended-phase testing are required under Marine Protection, Research, and Sanctuaries Act (MPRSA) Section 103 to determine the suitability of the material to be dredged for ocean disposal. Section 103 requires that all proposed operations involving the transportation and discharge of dredged material into ocean waters be evaluated to determine the potential environmental impact of such activities. The proposed placement must be evaluated using criteria published by EPA in Title 40 of the *Code of Federal Regulations* (40 CFR), Parts 220–223. Specific testing methods are described in the *Evaluation of Dredged Material Proposed for Ocean Disposal—Testing Manual* (EPA and USACE 1991, referred to here as the 'Green Book') and the *Evaluation of Dredged Material Proposed for Discharge in Waters of the U.S.*—Testing Manual (Inland Testing Manual or ITM) (EPA and USACE 1998). In addition, the EPA Refiner guidance manual, *Guidance for Performing Tests on Dredged Material Proposed for Ocean Disposal* (RTM) (USACE and EPA 2016) provides regional guidance on procedures for the two when assessing the suitability of dredge material for ocean disposal in EFA Refiner 2

The testing manuals provide guidance to support the tiered testing procedure for evaluating compliance with the limiting permissible concentration (LPC) as defined by the ocean dumping regulations. The procedure includes levels of increasing intestigative intensity that provide information to make ocean disposal decisions and is comprehensive enough to enable sound decision-making without unnecessary expenditure of time and recources.

1.2.2 Objectives and Deliverables

Evaluation of proposed dredge material from the project area pursuant to MPRSA Section 103 is required for ocean disposal of dredged material. For this reason, USACE Jacksonville District contracted with ANAMAR Environmental consuling, Inc. to collect sediment samples and to conduct physical, chemical, and toxicological caluations as required in 40 CFR Parts 220–228 and outlined in the testing manuals mentioned above.

Throughout the course of this roject, the procedures and criteria set forth in the Sampling and Analysis Plan/Quality Assurance Project Plan (SAP/QAPP) for sediment characterization were followed (Appendix A, ANAM P 2020). The objectives of this effort were to

- Collect the required volume of representative sediment samples from selected stations within the project area and the reference station within positioning accuracy appropriate for the project objective
- Collect and contanerize sediment samples according to proper protocols to ensure sample integrity.
- Test and characterize sediment samples for physical characteristics and chemical obtainments of concern and to perform toxicology bioassays in accordance with the Green Book and the RTM to determine the suitability of the materials for ocean disposal.
 - Deconstrate environmental compliance of sediments to be dredged and obtain concurrence compliance for offshore disposal of dredged sediments from USACE and EPA according to requirements specified in the Green Book, ITM, and RTM.
 - Provide a report to USACE and EPA on behalf of USACE in the format outlined in Section 6.2.6 of the SAP/QAPP (Appendix A).



Deliverables for this project include:

- An MPRSA Section 103 sediment testing report (draft and final) and supporting documentation that describe all aspects of the study and present the results of field sampling, physical and chemical analysis of sediment samples, and toxicological bioassays. This report presents comparisons of test sediments to the reference and provides the basis for a scientific recommendation regarding the acceptability of the dredged material for ocean disposal Important components of this report include:
 - A narrative addressing all aspects of field sampling, laboratory analysis, discussion of laboratory results, and a review of all laboratory quality control measures.
 - Laboratory results provided in condensed tables.
 - Maps of the sampling sites.
 - A Chemical Quality Assurance Report (CQAR [Appendix E]), which evaluates all representative data from the project field sampling and laboratory analyses. The CQAR summarizes the overall usability of the data for its intended surpose.
 - Daily Quality Control Reports (DQCRs) (Appendix B) prepared by the project manager for each day of field sampling.

ANAMAR coordinated and directed operations for this project and worked closely with USACE and EPA to develop sampling and analysis schemes, schedules and deliverables. ANAMAR also reviewed all data and produced this report summaring the esults of the physical, chemical, and toxicological analyses of sediment samples collected from the project area. Exhibits 1-3 and 1-4 indicate the principal data users and their respective areas of responsibility and subcontractors associated with this evaluation.

	Area(s) of Responsibility
USACE Jacksonville, FL	Responsible for maintenance and harbor improvements in the San Juan Harbor Federal Navigation Project and co-managing the San Juan Harbor ODMDS with EPA Region 2
EPA Fegion 2 Net York, NY	Give concurrence to environmental requirements of dredged sediment for approval for offshore disposal per the Green Book (EPA 1991), the ITM (EPA 1998), and <i>Guidance for</i> <i>Performing Tests on Dredged Material Proposed for Ocean</i> <i>Disposal</i> (USACE and EPA 2016)

		· /		
Evhibit 1_2	Principal Data Hears the	ЧП	N CICIO	Makore Accordant with This Project
	Fincipal Data Users a	5	ECISIC	Makers Associated with This Project



Exhibit 1-4. Subcontractors and Responsibilities Associated with This Report

Company and Contact Information	Area(s) of Responsibility
<u>Vibracore Subcontractor: Athena Technologies, Inc.</u> Project Manager: Adam Freeze P.O. Box 68, McClellanville, SC 29458 Phone: (843) 887-3800 <u>adam_freeze@athenatechnologies.com</u>	Vibracore support for field sample collection
Chemistry Laboratory: Analytical Resources, Inc. (ARI) Project Manager: Kelly Bottem 4611 S. 134th Pl., Ste. 100; Tukwila, WA 98168-3240 Phone: (206) 695-6211 kelly.bottem@arilabs.com	Laboratory sample preparation and chemical analysis of sediment elutriate, and tissues; sample hilding and archiving
Chemistry Laboratory: Materials Testing Consultants (MTC) Project Manager: Beth Goble 2118 Black Lake Blvd SW; Olympia, WA 98512 Phone: (206) 241-1974 beth.goble@mtc-inc.net	Preparation of electricate samples
Chemistry Laboratory: ALS Environmental, Inc. Project Manager: Todd Poyfair 1317 S. 13th Ave; Kelso, WA 98626 Phone: (800) 577-7222 Todd.Poyfair@alsglobal.com	Analysis of metals in the elutriate and situation samples
Geotechnical Laboratory: Terracon Project Manager: Chris Martin, Sr. 8001 Baymeadows Way Jacksonville, FL 32256 Phone: (904) 900-6494 crmartin2@terracon.com	Laboratory sample preparation and physical analysis of sediment; sample holding and archiving
Toxicology Laboratory: EcoAnalysts Project Manager: Brian Hester 4729 NE View Drive, Port Gambo, WA 98364 Phone: (360) 297-6040 bhester@ecoanalysts.com	Laboratory sample preparation and analysis for suspended phase, solid phase, and bioaccumulation potential
<u>Offshore Vessel</u> J.A.W. Marine Contractors, Inc. <i>Kruger B</i> Resularch Vessel San Juan, Peorto Rico	Support for field collection of sediment and water samples from the designated offshore reference station



2 MATERIALS AND METHODS

2.1 Project Design and Rationale

Areas proposed to be dredged were divided into five sampling areas representing associated dredging units or reaches (see Exhibits 1-2 and 2-1). All sampling stations were selected by USACE and approved by EPA Region 2. EPA reviewed available geotechnical data for borings taken in the areas to identify sediment strata horizons for informing the sampling and analysis plan. According to the scope of work, stiff clay was observed at depths of -34 to -35 fee MLW along the margins of Army Terminal Channel, between depths of -40- to -42 feet MLW when th Army Terminal Channel, and between depths of -45 to -46 feet MLW within the Anegado Chancel. Therefore, material associated with deepening below the channels themselves (currently at -40 feet MLW plus overdepth) is expected to be composed of stiff clay. In San Alcone Channel, sediment transitioned from gray silts and clays to sands and sand/clays at approximately -34 to -35 feet MLW. Existing channel depths in this area are -30 feet MLW plus overdepth. A brief description of each dredging unit is provided below.

<u>Reach A</u>: Composed of one composite of maintenance material (i.e., Mantenance M-A) collected from above the sand/clay interface from 10 locations spanning the Fastern Cruise Basin, Anegado Channel, Graving Dock, Puerto Nuevo and Army Terminal channels and Turning Basins.

<u>Reach B</u>: Composed of one composite of maintenance material (i.e., Maintenance M-B) collected from above the sand/clay interface from six locations spanning the Western Cruise Basin, San Antonio Channel, and San Antonio Approach Channel.

<u>Army Terminal Widener</u>: Composed of one composite of widening/deepening material (i.e., D-ATw-S) collected from above the clay interface (or project depth) from four stations in the widening area along the Army Terminal Channel

San Antonio Extension: Composed of one composite of widening/deepening material (i.e., D-SAx-S) collected from above the clay operative from three stations in the San Antonio Extension.

<u>Deepening</u>: Composed of one composite of deepening material (i.e., D-B-D) collected from below the sand/clay interface from to tomine stations in San Antonio Channel, San Antonio Approach Channel, San Antonio Exercice, and Western Cruise Basin.

<u>Clay Samples:</u> Completed condividual dense clay samples collected bottoms of cores, where encountered from sortions in Anegado Channel (D-AC-C), Eastern Cruise Basin (D-EC-C), Army Terminal Channel/Tuning Basin (D-AT-C), and Army Terminal Widener (D-ATw-C). Individual samples were inalyzed for physical and sediment chemistry parameters only.

<u>Reference</u>: The reference sediment (SJH20-REF) was collected from an offshore area in the vicinity of the San Juan ODMDS that has not been impacted by dredged material disposal. The reference station location was selected by EPA and is the same location that was sampled during the 2.15 MPRSA Section 103 evaluation for San Juan Harbor.

Analyses of composite samples consisted of three analytical tiers, including sediment physical and chemical (sediment, elutriate, and tissue) analyses and toxicological bioassays. A summary of field sampling methods used during the collection process are presented in Exhibit 2-2. Sediment samples were analyzed for the contaminants of interest and bioassay test species listed in Exhibit 2-3.



Dredging Unit/Reach	Subsample IDs	Estimated Mudline Elevation (ft, MLLW) ^[2]	Project Elevation Including 2' Allowable Overdepth (feet MLLW) ^[1]	Est. Core Length Project Lepth ^[2]	Notes
	M-A-S-1	-42.5	mudline to -48		Yellow highlight
	D-AC-C-1 (clay)	-42.5	mudime to -46		indicates sediment elevations below
	M-A-S-2	-31.6	mudline to -37	5.4	target project depth.
	D-EC-C-1 (clay)				Maintenance (surface)
	M-A-S-3	-30.7	mudline to -37	6.3	material is considered the unconsolidated
	D-EC-C-2 (clay)				layer of material above
	M-A-S-4	-37.4	mudline o -3	-1.4	the native material.
M-A	M-A-S-5	-39.2	mut the to	-0.2	Deepening (clay)
(SJH Maintenance Reach A)	M-A-S-6	-38.7	mudline -39	0.3	material is considered native material.
	M-A-S-7	-40.5	mudline to -39	-1.5	
	M-A-S-8	-40.9	mudline to -46	4.2	
	D-AT-C-1 (clay)				
	M-A-S-9	-34.4	mudline to -34	-0.4	
	D-AT-C-2 (clay)				
	M-A-S-10		mudline to -46	4.8	
	D-AT-C-3 (clay)	10.0-		ч.0	
	D-ATw-S-1	-19	mudline to -44	25.0	Maintenance (surface material is considered
	D-ATw-C-1 (clay)	V			the unconsolidated layer of material above
Army Terminal Widener	D-A1 w-S-2 D-A1 C-2 (clay	-16	mudline to -44	28.0	the native material.
	D-ATLS-3	-19.4	mudline to -44	24.6	Deepening (clay)
	D-ATw-S-4 D-Atiw-C-3 (clay)	-16	mudline to -44	28.0	material is considered native material.
K.			8		

Exhibit 2-1. Summary of Sampling Scheme Including Dredging Units, Elevations, and Estimated Care Lengths

MPRSA Section 103 Sediment Characterization San Juan Harbor, Puerto Rico



Dredging Unit/Reach	Subsample IDs	Estimated Mudline Elevation (ft, MLLW) ^[2]	Project Elevation Including 2' Allowable Overdepth (feet MLLW) ^[1]	Est. Core Lengtl to Project Depth (feet)	Notes
	M-B-S-1				Yellow highlight indicates sediment
	D-B-D-1	-30.7	mudline to -38		elevations below target project depth.
	M-B-S-2	07.7	III. (00 ♦	0	Maintenance (surface)
	D-B-D-2	-37.7	mudline to -38	0.3	material is considered the unconsolidated
	M-B-S-3	24.2			layer of material above the native material.
M-B	D-B-D-3	-34.2	mudline to 38	3.0	Deepening material is
(SJH Maintenance Reach B)	M-B-S-4	20.4	V2		considered native material.
	D-B-D-4	-38.4	mudit e to -38	-0.4	material.
	M-B-S-5	-35.5	udline to -38	0.5	
	D-B-D-5			2.5	
	M-B-S-6	-43.9	mudline to -38	-5.9	
	D-B-D-6				
	D-SAx-S-1	X	mudling to 20	0.5	Yellow highlight indicates sediment
	D-B-D-7	38.5	mudline to -38	0.5	elevations below target project depth.
	D-SAx-S-2	O _{-38.0}	mudling to 20	0.0	Maintenance (surface)
San Antonio Extension	D -D-8	-38.0	mudline to -38	0.0	material is considered the unconsolidated
	D-SA-S-3				layer of material above the native material.
	D-B-D-9	-28.4	mudline to -38	9.6	Deepening material is considered native
					material.
K K	0				
	▼				
			9		

MPRSA Section 103 Sediment Characterization San Juan Harbor, Puerto Rico



Dredging Unit/Reach	Subsample IDs	Estimated Mudline Elevation (ft, MLLW) ^[2]	Project Elevation Including 2' Allowable Overdepth (feet MLLW) ^[1]	Est. Core Lengtl to Project Depth (feet)	Notes
SJH20-REF (Reference station)	N/A	N/A	Grab sample	W A	
Site Water Stations					
SHJ20-SW	N/A	N/A	Collect 1 m above bottom	N/A	
SJH20-REF-SW	N/A	N/A	Collect 1 m below surfac	N/A	

[1] Project elevation is the authorized deepening depth plus allowable overdepth below MLLW (feet

[2] Mudline elevation and estimated core length based on August 2020 bathymetric survey da

 \mathbf{X} -XX-X-X = maintenance (M) or deepening/widening material (D)

X-X-X = San Juan Harbor Maintenance Areas (A and B); Army Terminal (AT); Anegada Channer (AC); San Antonio (SA); Army Terminal widening (ATw); San Antonio extension (SAx)

X-XX- \mathbf{X} -X = surface stratum (S), deep stratum (D), or clay (C)

X-XX-X-X =station number

MLLW = mean lower low water



Exhibit 2-2. Summary of Field Sampling Materials and Methods

FIELD SAMPLE COLLECTION:

• Project sub-samples and composite samples from each dredging unit plus reference sediment

SAMPLING GEAR:

- Project samples collected by vibracore or grab sampler
- Reference sediment collected with double van Veen sampler
- Water parameters measured with YSI multiprobe meter and Hach 2100P turbidimeter

PRESERVATION:

- Sediment samples were kept at or below 4°C
- Holding-time requirements were analyte-specific and test-specific

IN SITU WATER COLUMN DATA:

Conductivity (mS/cm) pH Sea state Turbidity (NTU) Water temperature (°C) Dissolved oxygen (mg// anc /o saturation) Salinity (ppt) Tide cycle Water depth (feet) Weather observations

Exhibit 2-3. Analytical Requirements Per Sample Collected

	Sample:			Clay				Pre-exposure
Test			Subsamples	Sample	Reference	Control	Site Water	Tissues
	Grain Size	Y	Y	Y	Y	Y		
6	Atterberg Limits	Y			Y			
Physicals	% Moisture	Y	Y	Y	Y	Y		
hys	Settling Rates	Y	🗙		Y			
	Specific Gravity	Y			Y			
	Bulk Density	Y			Y			
	TOC	Y		Y	Y	Y		
ent stry	Metals	Y		Y	Y			
Sediment Chemistry	Pesticides			Y	Y			
S S	PCB Congeners	6 Y /		Y	Y			
	PAHs	Y		Y	Y			
e Ly	Metals	Y					Y	
Elutriate Chemistry	Pesticia	Ý					Y	
Elu Che	PCB Congeners	Y					Y	
	M. tals	Y			Y			Y
le stry	Pesti des	Y			Y			Y
Tissue	P/Congeners	Y			Y			Y
	AH	Y		-	Y			Y
	ipids	Y			Y			Y
λβ	Suspended Phase Bioassay	Y				Y		
Toxicology	Solid Phase Bioassay	Y			Y	Y		
Τc	Bioaccumulation Potential	Y			Y	Y		

Y = analysis performed; -- = analysis not performed/not required or not applicable

PCB = polychlorinated biphenyl; PAH = polycyclic aromatic hydrocarbon



2.2 Sample Collection Techniques

2.2.1 Project Field Effort

Sampling activities were conducted according to the SAP/QAPP (Appendix A) and guidance from USACE and EPA. Field mobilization and sampling took place from October 12 through November 2, 2020. Field personnel consisted of scientists from ANAMAR and Athena Technologies. The *Kruger B* vessel departed from Pier 9 of the Port of San Juan for collection of the reference sediment and water on October 29, 2020. The Athena vessel *Good Vibrations* was used to collect the project samples and site water within the project area. Sample composition was conducted on-site by ANAMAR personnel prior to shipping samples to the laboratorics.

Exhibit 2-4 is a summary of the field sampling, compositing, and shipping activities. For more details, refer to the DQCRs in Appendix B. Breaks in the field sampling structule reflect mobilization and collection of samples at additional project sites.

Exhibit 2-4. Field Sampling Activities

Date	General Activity
Oct 12 and 19, 2020	 Mobilize to San Juan, PR; get boat out of ustoms and stage equipment to begin sampling operations
Oct 19, 2020	Begin collection of sediment sampler from Peach A
Oct 20, 2020	 Finish collection of sediment samples, on keach A
Oct 21, 2020	 Begin collection of sediment samples from Army Terminal Widener Begin compositing samples
Oct 22, 2020	 Finish collection of sediment samples from Army Terminal Widener Start collection of sediment samples from Reach B and San Antonio Extension Continue compatition samples
Oct 23, 2020	 Finish collection of seament samples from Reach B and San Antonio Extension Fini in compositing samples Begromaking arrangements for shipment of samples
Oct 26, 2020	 Cellect te water and background water chemistry kit
Oct 27, 2020	 Packing prepare project sediment and water samples for shipping Prepare chains of custody Sky samples to laboratories via FedEx Custom Critical
Oct 29, 2020	Collect offshore reference sample and watersample
Nov 2, 2020	 Pack and prepare reference sample for shipping Prepare chains of custody Ship samples to laboratories via FedEx Custom Critical

2.2.2 Site Positioning

Sediment sampling locations were provided by USACE and approved by EPA. Station correctes were uploaded to a Panasonic Toughbook computer and associated Trimble submeter GPS system on the R/V *Good Vibrations* and a GPS system at the helm of the S/V *Kruger B*. A Garmin hand-held GPS was used to log sampling coordinates at the aft deck of the *Kruger B* during sampling. Sampling coordinates were also logged at coring stations with a Garmin hand-held GPS as back-up. Waypoints were recorded on sampling field logs. Navigation and positioning of the sampling vessels referenced above were handled by U.S. Coast Guard-licensed captains under direction of the ANAMAR field team leader. A graduated line was used



to determine water depths at coring locations. Water depths during offshore grab sampling were determined using a depth finder.

All samples were taken within 50 feet of the target station and conformed to Subsection 11.1.3 of the SAP/QAPP. Table 1 contains dates and times, coordinates, water depths, bottom elevations, and associated data for sediment grab and core samples. Table 2 contains similar information for water column parameters recorded at the reference station and the site water location within San Juan Harbor. The sampling locations for reference and project sediment samples are shown in Maps 1 through 5.

2.2.3 Decontamination Procedures

All equipment contacting sediment samples was cleaned and decontaminated as described below. Work surfaces on the sampling vessel were cleaned before the sampling valuegan and before leaving each station. All equipment contacting sediment samples was depontaminated between dredging units and individual stations, where required, to prevent coss-contamination. Gloves used at a given sampling station were replaced with new gloves nor to sampling at the next station.

Decontamination Procedures

- · Wash and scrub using site water or tap water to remove gross contamination
- Wash and scrub with Liquinox detergent
- Rinse with site water
- Rinse with deionized water
- Rinse 2 times with pesticide grade isopropagilation
- Rinse 3 times with deionized wher
- Equipment not being used immediately was air-dried and stored wrapped in new, clean aluminum foil

Any derived waste was contributed and disposed of in accordance with federal, state, and local laws.

2.2.4 Water Column de succements

A YSI multiprobe mean and a Hach 2100P turbidimeter were used to measure water column parameters at the defended site water station and at the San Juan Harbor site water station. Instruments were can rated each day prior to use according to manufacturer's instructions. An end-of-day reading was also taken to document that the instrument remained calibrated within acceptance criterion. Water column measurements were recorded from 2 or 3 feet below surface, at mid-dooth, and 3 feet above the bottom at the San Juan Harbor site water station. Water columneesurements were taken 2 feet below the surface at the reference station. Measured water, ohim parameters and associated data consisted of

- Time of reading
- Depth of measurement (feet)
- Water temperature (°C)
- pH (units)
- Salinity (parts per thousand [ppt])
- Conductivity (mS/cm)
- Dissolved oxygen (mg/L and percent saturation)
- Turbidity (NTU, near-surface only)



Water depth measurements, tidal cycle, and weather observations were recorded on field logs and are summarized in Table 2. Equipment calibration logs are in Appendix B.

2.2.5 Sediment Sampling with Vibracore

Subsurface core samples were obtained using a vibratory core sampler (vibracore). Vibracore services were performed by Athena Technologies under the guidance of an ANAMAR field teap leader who was present on the sampling vessel at all times to direct operations, record field press, and containerize and label samples. The vibracore samples were collected from the sampling vessel *Good Vibrations*, which is fitted for vibracore sampling. The vessel carried all necessary sediment sampling equipment and materials.

The vessel captain navigated to each target using a helms map displayed anasonic Toughbook computer and associated Trimble GPS system. Once on-station, vessel was immobilized using a three-point anchoring system. Vessel coordinates w oppared to station re coordinates loaded in a second GPS to confirm location accuracy. Dept recorded to the wer nearest inch using lead-line readings and were then converted to the nearest tenths of a foot. Bottom elevation was calculated in the field using real-time water level tata (feet MLLW) from National Oceanic and Atmospheric Administration [NOAA] Station 55371 at San Juan. Core penetration required to reach project depth was calculated by dding real-time elevation of the substrate surface (as a negative value) to the project depth

Athena's vibracore system was deployed from the deck of the vessel and consisted of a generator with a mechanical vibrator attached via cable. This vibrator was attached directly to a 4-inchdiameter stainless steel core barrel. The sampler was lowered to the substrate through a moon pool in the deck of the sampling platform by statisting lengths of drill stem. The vibracore apparatus was then activated and the correbarred penetrated into the sediment until it reached target depth or refusal, whichever was mach penetration into the sediment at the vibracore drill stem does not result in mean table penetration into the sediment. This is often the result of the end of the coring tube end untering rock or consolidated sediment.

When the vibracore reached taget depth or refusal, the vibracore apparatus was deactivated and the core retrieved using an electric winch. Once the sample was on-deck, the recovered core length was determined to the nearest inch and converted to the nearest tenths of a foot. Determination of a spin one of a given core sample was based on percent recovery requirements as stated in the SAPDAPP. The sediment sample was then removed from the core barrel and placed into a spinles steel bin for characterization, photographs, and containerizing.

When soliment cores are collected with a vibracore system, the retrieved sample is subject to match compaction. For instance, a core sample taken from a penetration depth of 10 feet may result in precovered core of only 8 to 9 feet in length, depending on the sediment composition. Core samples were considered acceptable if the core was inserted vertically into the sediment, reached target depth or refusal, and recovered at least 75% of penetrated depth. Alternatively, the acceptance limit for each core was decreased if the first core attempted at a given station was below 75% recovery of penetration depth and subsequent cores collected were within ±15% of the initial core percent recovery. During events when collected cores showed widely varying recoveries over several attempts, the material was collected, and the recovery lengths and reason for low recoveries were recorded on the field sheets.

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The number of cores collected at each station was dictated by the number needed to achieve sufficient volume for laboratory analyses. To maintain proportional volumes between subsample stations, the team tried to collect the same number of cores at each station. However, in some circumstances, it was difficult to predict how many cores would be required at each station across a dredging unit because of the requirement to separate out the clay layer from the overlying unconsolidated material. Also, some stations within a dredging unit had less than 2 feet of shoaling and therefore required a grab sample. EPA was consulted on this issue and it was recommended that if an equal number of cores could not be collected, then an equal volume o material should be collected at each station.

Once all cores or grab samples were collected at a given station, the sample material cas photographed, transferred to labeled Teflon[®] bags, and placed into ice-filled coolers. All containers were properly labeled, and sampling information for each station was recorded on individual project-specific field logs. At the end of each sampling day and followin, compositing, the samples were transferred to a refrigerated truck for storage at $\leq 4^{\circ}$ C proves shipping.

Information from core logs is summarized in Table 1. Field sampling logs are in Appendix B. Photographs taken during sampling and compositing are in Appendix *I*.

2.2.6 Sediment Sampling with Grab Sampler

Within the project reaches, there were some stations with very short cores lengths or areas where the mudline elevation was below the project depth. BPA actived that if shoaling was <2 feet above the target project depth or the mudline elevation we below project depth, a grab sampler could be used to collect the material. Grab samples were collected using either a double van Veen (for the reference station) or a modified Petersen grab sampler that was lowered and raised by a winch. One person operated the wince and additional team members guided the sampler into a decontaminated stainless steel in or the vissel. Excess water was allowed to drain from the sampler prior to placing sample many all the bin. When the required volume of sediment was collected, a photograph of the materia was taken and notes on the sample's appearance and characteristics were recorded on a project-specific field log. Using decontaminated stainless steel utensils and disposable buile goves, the sample was placed in pre-cleaned, labeled Teflon[®] bags and stored in ice-filled coorts. Upon return to the dock, the samples were transferred to a refrigerated truck for preserv ion at or below 4°C. Map 1 shows the location of the reference in Appendix B provide additional information on grab sampling. station. Table 1 and th lo Photographs taken d pling and compositing are in Appendix I. ing s

2.2.7 San ble Processing, Shipping, and Custody

2.2.7.1 Com ositing and Homogenizing

ANAMA2 personnel composited and homogenized sediment samples using decontaminated stainless seel mixing equipment and a 40-gallon-capacity stainless steel bin. Compositing was conducted in accordance with the scheme presented in Section 2.1 with the following exception. No DB-II composite sample was collected. None of subsample stations within the D-B-D dialogue unit had material representative of deepening or native sediments above the project deput; therefore, no deepening samples were collected. See Table 1 for more information.

After sediment samples were composited, appropriate volumes of each sample were divided and placed in method-specific, pre-cleaned, pre-labeled Teflon[®] bags or glass jars (for chemical analysis) or plastic bags or buckets (for physical analysis or for use in bioassay testing). Once composited, the samples were placed in a refrigerated truck at or below 4°C until shipment to



respective laboratories. The temperature inside the truck was monitored to ensure that samples met preservation criteria. Copies of temperature logs are in Appendix B.

2.2.7.2 Shipping to Laboratories

Samples were placed in refrigerated units called C-Safes and shipped to laboratories overnight via FedEx Custom Critical. The temperature within the C-Safes was monitored throughout the shipment. Copies of temperature logs are in Appendix B.

Chain-of-custody records for each laboratory were completed to reflect the final sample name and to identify the analyses and analytical methods required. These chain-of-custody form accompanied the samples during shipment to the laboratories. Copies of final signed chain-ofcustody forms are included with the laboratory reports in Appendices C, D, and E

2.3 Physical and Chemical Analytical Procedures

2.3.1 Physical Procedures

Terracon performed physical analysis of all sediment samples. AN MAX performed quality assurance/quality control (QA/QC) on sediment physical data and presented the data for all samples in summary tables.

2.3.1.1 Grain Size Distribution

Gradation tests were performed in accordance with methods ASTM D-422 and ASTM D-1140. Each representative sample was air-dried and dry-prepped in accordance with method ASTM D-421, and results of the sieve analysis of material larger than a #10 sieve (2.00-mm mesh size) were determined. The minus #10 sieve material was then soaked in a dispersing agent. Following the soaking period, the sample was placed in a mechanical stirring apparatus and then in a sedimentation cylinder where hydroteeur realings were taken over a 24-hour period. After the final hydrometer reading was taken, we sample was washed over a #200 sieve (0.075-mm mesh size), placed in an oven and dried to a constant weight. After drying, the sample was sieved over a nest of sieves to determine the gradation of the material greater than #200 sieve size. Cumulative frequency proceedees were graphed and presented by Terracon on USACE Form 2087 (Appendix C).

2.3.1.2 Moisture Contern

Moisture content has performed in accordance with method ASTM D-2216-80 and Plumb (1981). The sample weight was accorded and the sample was placed in an oven and dried to a constant mass at 110°C. Once a constant dry mass was obtained, the percent moisture was determined by subtracting the dry mass from the wet mass, then dividing the loss in mass due to drying (the mass objust moisture) by the wet mass. The percent total solids was reported on a 100% wet weight basis.

Atterberg Limits

Texts of liquid and plastic limits were performed in accordance with ASTM D-4318, wet method, as follows. The minus #40 sieved material was mixed with a small amount of water and placed is a liquid limit device. A groove was cut using a flat grooving tool and the liquid limit was determined by the number of drops of the cup. When the number of drops was in the desired range, a moisture sample was obtained, placed in a 230°C oven, and dried to a constant weight. This was repeated until three determinations had been obtained, one between 15 and 25 blows,



one between 20 and 30 blows, and one between 25 and 35 blows. The reported value is the intersecting value at 25 blows when all three are plotted.

The plastic limit was determined by slowly air-drying a small sample left over from the liquid limit determination. The sample was rolled and air-dried until the thread became crumbly and lacked cohesion. When this point was reached, the sample was placed in a tare and weighed, then placed in an oven and dried to a constant weight. The moisture content is the plastic limit.

2.3.1.4 Specific Gravity

Specific gravity was determined in accordance with method ASTM D-854. Each sample was placed in a mechanical stirring device and deionized water was added to form a slurdy. The sturry was then transferred to a pycnometer and was de-aired by applying a vacuum. After vacuuming, the pycnometer with sample was allowed to reach thermal equilibrium. The vaculevel was adjusted to a calibration mark, and the pycnometer with sample was weighted. After the pycnometer with sample weight was recorded, the sample was emptied into a drying container and placed in an oven until a constant dry mass of sediment solids was cotained.

2.3.1.5 Bulk Density

Bulk density, also known as dry bulk density, is the weight of dry segment divided by the total volume. The total sediment volume is the combined volume of solids and pores which may contain air, water, or both. The average values of air, water, and solids in soil are easily measured and are a useful indication of the sediment's physical condition.

2.3.2 Chemical Analytical Procedures

ARI and ALS performed chemical analyses of the sample composites and the reference in accordance with published procedures. Analytical methods, preparation methods, target detection limits, and laboratory reporting noits for sediment are in Subsection 13.3 of the SAP/QAPP (Appendix A). ANAMAR penermed QA/QC on these data and presented them in summary tables. Complete laboratory reports are in Appendix D. Exhibit 2-5 provides a summary of analytical methods for chemical analysis of sediment.



Exhibit 2-5.	Summary of Methods and Equipment Used during Chemical Analysis of
	Sediment

5eai	ment	
EPA Method	Instrument/ Procedure	Methodology Summary
6020A (Trace metals in water/sediments/ tissues)	Inductively Coupled Plasma/Mass Spectrometry	Inductively coupled plasma/mass spectrometry (ICP/MS) is applicable to the determination of sub-µg/L concentrations of a large number of elements in water and sediment samples. Acid digestion prior to filtration and analysis is required for aqueous samples and sediments for which total (acid-leachable) element are required. For analysis, sample material in solution is introduced by pneumatic nebulization into radiofrequency preme where energy transfer processes cause desolvation atomization, and ionization. The ions are extracted from the clasma through a differentially pumped vacuum interface and separate for the basis of their mass-to-charge ratio by a quadra alternass spectrometer. The ions transmitted through the dustrupole are detected by an electron multiplier and the ion information is processed by a data-handling system.
7470/7471 (Mercury in water/sediments/ tissues)	Mercury Analyzer Cold Vapor Atomic Absorption	Method 7470 is applicable to wate samples, and 7471 is applicable for measuring total mercury (organic and inorganic) in sediments. All samples are digest than oxidized at $95 \pm 3^{\circ}$ C, then mercury from the digest tes is reduced to the elemental state and aerated from solution a closed system. The mercury vapor passes through a coll resitioned in the light path of an atomic absorption spectrophotometer, and the absorbance (peak area) at 253 Jum is measured as a function of mercury concentratio
8081/8082 (Pesticides/PCBs in water/ sediments/ tissues)	Gas Chromatoghtah	Method 1981 and 8082 are applicable to the determination of xtrailed or anochlorine pesticide compounds and typic or an additional displayed by the second strain of the secon
8270 (CAHSin settimen tissu (s)	Gas Chromatograph/ Mass Spectrometer	This method is used to determine the concentration of polycyclic aromatic hydrocarbon (PAH) organic compounds in extracts prepared from many types of solid matrices and water samples. The extracted sample aliquot is injected into a gas chromatograph/mass spectrometer (GC/MS) by large-volume injection for qualitative and quantitative determination. Data may be obtained from the mass spectrometer via one of the three modes of operation: full scan mode, selected ion monitoring (SIM), or multiple reaction monitoring (MRM).
Plumb (1981) (TOC in sediments)	Total Organic Carbon (TOC) Analyzer	Plumb (1981) is used to determine the concentration of organic carbon in sediment by catalytic combustion or wet chemical oxidation. The carbon dioxide formed from this procedure is measured and is proportional to the TOC in the sample.



2.4 Bioaccumulation and Toxicology Procedures

EcoAnalysts conducted biological testing using sediment samples collected by ANAMAR as part of the dredged material evaluation for San Juan Harbor. The testing procedures used by EcoAnalysts (2021) is summarized in Section 2 of their report titled *Toxicity Testing Results, San Juan Harbor Puerto Rico 103 Evaluation, San Juan, Puerto Rico*. The complete laboratory report is in Appendix G (in hardcopy and on disc).

The material under consideration for ocean disposal was evaluated in accordance with procedures and criteria outlined in the Green Book and the RTM and with guidance outlined in the ITM. Biological analyses with reference sediments was performed concurrently with the text sediment evaluations.

This program included bioassay analysis of four composite samples and one reference sample. In addition, appropriate laboratory control samples (LCSs) were run with each of us selected test species. Bioassay testing for this project consisted of three water column bioassays, two whole sediment bioaccumulation potential tests. The bioassay and bioaccumulation tests are summarized in Exhibit 2-6. Exhibit 1-7 summarizes the testing objectives for each sample evaluated under this program. All tests were conducted within the eight-week (56 days) sediment holding time limit.

Test Type	Type of Organism	Тахо	Project Sediments	Reference Sediment	Control Sediment/ Seawater
	Mysid shrimi	Amerikamysis buhia Courtesy of: Alan Kennedy, ERDC	•1	NA	•
Suspendud- Particulate Phale	Fish	Menidia beryllina	•1	NA	•
	Larval bivalve	Mytilus galloprovincialis	•1	NA	•

Exhibit 2-6. Biological Testing Performed for Dredge Material Evaluation

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Test Type	Type of Organism	Taxon	Project Sediments	Reference Sediment	Control Sediment/ Seawater	
Solid-Phase	Amphipod	Ampelisca abdita	•		Ó	
	Mysid shrimp	Americamysis bahia	j.	(C)	•	
Bioaccumulation	Polychaete	Alitta virens		•	•	
	Bive	Macona nasuta	•	•	•	

¹ Sediment elutrices of project material NA = Tests or treaments that are not applicable to the selected tests.

Exhibit 2-7. Biological Testing Objectives by Sample

its



2.5 **Tissue Analysis Recommendations**

ANAMAR coordinated with USACE and EPA to determine which analytes should be tested in the corresponding tissue samples based on guidance provided in the RTM. The final list of parameters analyzed in tissue samples is summarized in Exhibit 2-3.

2.6 Applicable Technical Quality Standards

Raw field and laboratory data were summarized and compiled into tables. Figures were used t associate the results spatially with respect to sampling locations.

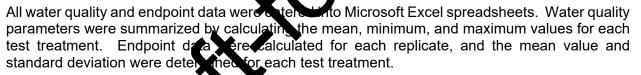
2.6.1 Sediment Chemistry

Results of laboratory analyses of sediment samples are compared to published sediment screening values as appropriate and in conformance with the Green Book and the ACM. These levels are the threshold effects level (TEL) and effects range low (ERL). The TELE presents the concentration below which adverse effects are expected to occur only are y. The ERL is the value at which toxicity may begin to be observed in sensitive species (Bochman 2008). These comparisons are for reference use only and are not intended for regulatory decision-making.

2.6.2 Elutriate and Water Chemistry

Analytical results for elutriate and water samples were compared to the latest published EPA water quality criteria of criteria maximum concentration CMC [synonymous with 'acute']) established in EPA (2006, 2015). The CMC is an estimate of the highest concentration of a pollutant in saltwater to which an aquatic community can be exposed briefly without resulting in an unacceptable effect (EPA 2006, Buchman 2008).

2.6.3 Toxicology



All hand-entered data were reviewed for data entry errors. Any errors found were corrected before summary calculations were performed. A minimum of 10% of all calculations and data sorting were reviewed for errors. For ew counts were conducted on any apparent outliers.

Statistical comparisons were made according to the Green Book and were performed using SAS/STAT solware or CETIS[™] software (CETIS 2012). Before statistical comparisons were conducted, data were tested for normal distribution. Any data that violated the assumption of normal distribution were transformed using an arcsine square root transformation before statistical analyses. Al data were tested for equality of variance using Levene's test.

Pentilic test results were compared to reference results using analysis of variance (ANOVA) with SAC Proc GLM software with Dunnett's multiple comparison test on the mean values. The Dunnett's test was performed as a one-way test, testing for significantly lower organism survival than in the reference sample.



2.6.4 Tissue Chemistry

The project sample and reference tissues had five replicates per test species and were evaluated using guidance from Subsection 6.3 of the Green Book and Subsection 9.2.3 of the RTM. Analytical results for tissue samples were compared to published tissue screening benchmarks. The U.S. Food and Drug Administration (FDA) action levels and threshold levels were used for comparison after accounting for steady-state adjustments as applicable.

Analyte concentrations in *Macoma nasuta* tissues were compared to FDA levels for birdve mollusks. Analyte concentrations in *Alitta virens* tissues were compared to the FDA levels for crustacea as there are no FDA levels published for polychaete worm tissue (FDA 2001, 2 11).

The mean of results for each set of five replicates per composite and analyte combination was calculated for wet weight and dry weight concentrations. The wet weight concentrations of composites having two or more replicates greater than the MDL were compared to the replicate concentrations for the reference tissue per analyte. Mean values of analyte concentrations were calculated as follows:

- For non-detects (U-qualified) data, the method detection (mit MDL) was used in all statistical calculations.
- For J-qualified and non-qualified analytical results are reported result was used in all statistical calculations.

In cases where the mean concentration of an analyte in *virens* or *M. nasuta* tissue was found to exceed that of the reference tissue, the biotatistical software program ToxCalc v5.0.32 (Tidepool Scientific, LLC) was used to determine the relative distribution and variances among the samples tested. If the distribution was determined to be abnormal or the variances unequal, the data were treated with a reciprocal random on and the distribution and variances were reevaluated. If no mean tissue contamined to concentration was found to statistically exceed that of the reference tissue, no additional analysis was necessary to demonstrate compliance with the LPC (Green Book). Mean tissue contained results found to statistically significantly exceed those of the reference tissue (of the came species) are presented in bold font in the accompanying tables. This is in accordance with Subjection 9.2.3 of the RTM.

2.7 Reporting Limits

MDE, and method reporting limit (MRL) were reported on a dry weight Chemical conc atic mes, liquid basis for site water and elutriate samples, and wet and dry basis for seg ment weight base for tissues samples. The MDL refers to the minimum concentration of a given analyte and reported with a 99% confidence level that the analyte concentration is that can be me ure greate The procedures for determining MDLs is defined in 40 CFR Part 136 Appendix than zero. B for mo chemical analyses. The MRL refers to the minimum concentration at which the www.report analytical chemistry data with confidence in quantitative accuracy of a given lab Common laboratory procedures for defining an MRL include assigning it to a fixed dat above the MDL or by using the lowest calibration standard. MRLs are often adjusted by boratory for sample-specific parameters such as sample weight, percent solids, or dilution.



3 RESULTS AND DISCUSSION

3.1 Field Data and In Situ Measurements

3.1.1 Weather Conditions

Conditions during sampling at the offshore reference station and coring locations were favorable Weather conditions (including wind direction, wind speed, and sea state) at each station are not if on the field logs in Appendix B.

3.1.2 Water Column Data

Water column parameters were recorded at the offshore reference station (SJH20-FEF-SW) and at the site water location within the San Juan Harbor project area (SJH21-SV) and are summarized in Table 2. The water sampling field logs are in Appendix B.

3.1.3 Vibracore and Grab Sampling Data

A brief summary of sample collection activities within each dredging unit is provided below. EPA was consulted throughout the sampling effort. Key issues that were discussed are summarized in Subsection 4.1. Table 1 provides a summary of vibracore sampling data, including core depth, penetration, recovery length, and percent recovery. Copies of the core logs are in Appendix B.

San Juan Harbor Maintenance Reach A Summary:

M-A-S-1/D-AC-C-1. Project depth of -48 feet MLLW was reached at this station. Three core samples were collected in liners. No hard, stiff clay layer indicative of native (new work) material was encountered. EPA Region 2 inspected the pract cores and determined that the material throughout the profile was characteristic of manufenance material. Therefore, no clay sample (D-AC-C-1) was collected.

M-A-S-2/D-EC-C-1. Project depth of -37 net MLLW was reached at this station. One core was collected in a liner and a second ore was collected with the 4-inch unlined core barrel. There was no obvious stratification network maintenance and deepening sediment. No hard, stiff clay layer indicative of native (new work) material was encountered. The EPA Region 2 representative inspected the intact core and Netermined that the material throughout the profile was characteristic of maintenance instead. Therefore, no clay sample (D-EC-C-1) was collected.

M-A-S-3/D-ECCO-D Project depth of -37 feet MLLW was reached at this station. One core was collected in a liner and a second core was collected with the 4-inch unlined core barrel. There was not obvices stratication between the maintenance and deepening layer, but the material did get slightly stiffe toward the bottom 3 feet of the profile. That slight transition is where the sample was spin between the maintenance and deepening. Therefore, a both a surface (M-A-S-3) and a clay sempting (D-EC-C-2) were collected.

M-A-1-4 infough M-A-S-7. EPA was consulted about the lack of material above project depth at the second stations. The EPA Region 2 representative advised that if the shoaling was <2 feet, a grab sample could be collected. Therefore, these four stations were collected with a grab ampler. Equal volumes were collected at each station.

M-A-S-8/D-AT-C-1. Project depth of -46 feet MLLW was reached at this station. One core was collected in a liner and a second core was collected with the 4-inch unlined core barrel. There was no obvious stratification between maintenance and deepening. No hard, stiff clay layer



indicative of native (new work) material was encountered. Therefore, no clay sample (D-AT-C-1) was collected.

M-A-S-9/D-AT-C-2. This area is already below the deepening project depth. The vessel captain tried to relocate the station but could not find any shoals above the project depth. Therefore, no surface or clay sample was collected at this station.

M-A-S-10/D-AT-C-3. Project depth of -46 feet MLLW was reached at this station. Two cores were collected and retained. There was no obvious stratification between maintenance and deepening. No hard, stiff clay layer indicative of native (new work) material was encountered. Therefore, no clay sample (D-AT-C-3) was collected.

San Juan Harbor Maintenance Reach B Summary:

M-B-S-1/D-B-D-1. Project depth of -38 feet MLLW was reached at this station. T to cores were collected at this station. No deepening (native) material was encountered. Therefore, no deepening sample (D-B-D-1) was collected.

M-B-S-2/D-B-D-2. Sediment elevation (-38.5 feet) at this station is below project depth of -38 feet MLLW. EPA was consulted and advised ANAMAR to the a grab sampler to collect unconsolidated maintenance material at the surface. Therefore no deepening sample (D-B-D-2) was collected.

M-B-S-3/D-B-D-3. Project depth of -38 feet MLLW was mached at this station. Two cores were collected at this station. No deepening (native) material was encountered. Therefore, no deepening sample (D-B-D-3) was collected.

M-B-S-4/D-B-D-4. Sediment elevation -391 feet at this station is below project depth of -38 feet MLLW. EPA was consulted and a vision ANAMAR to use a grab sampler to collect unconsolidated maintenance material at the surface. Therefore, no deepening sample (D-B-D-4) was collected.

M-B-S-5/D-B-D-5. Sediment en vation 39.2 feet) at this station is below project depth of -38 feet MLLW. EPA was consumed and advised ANAMAR to use a grab sampler to collect unconsolidated maintenancempterial at the surface. Therefore, no deepening sample (D-B-D-5) was collected.

M-B-S-6/D-P.D-6. Sediment elevation (-44.6 feet) at this station is below project depth of -38 feet MLLW. EPT was consulted and advised ANAMAR to use a grab sampler to collect unconsolidated pair enance material at the surface. Therefore, no deepening sample (D-B-D-6) was connected.

Arry Terminal Widener Reach Summary:

D-ATM-SIMD-ATw-C-1. Refusal was encountered (-36 feet MLLW) above the project depth on 44 Set MLLW due to hard, stiff clay. One core was collected from this station, and it had material characteristic of both maintenance and new work (native) material. Therefore, both a urface (D-ATw-S-1) and a clay (D-ATw-C-1) sample were collected.

D-ATw-S-2/D-ATw-C-2. The length of core required to reach project depth of -44 feet MLLW was longer than could be reached with a 20-foot core barrel (target penetration = 27.9 feet). These limitations were discussed with EPA prior to sampling, and a "stair-step" method was suggested that involves collecting another core downslope of the target location to reach full project depth



(or refusal by encountering native material). This approach was required at this station because the core penetration length at the target location was 19.6 feet (bottom core elevation of -35.7 feet MLLW) but did not encounter refusal. Therefore, ANAMAR consulted with EPA while on station for approval to use the "stair-step" approach. A second station was located downslope of the target location with a top of core elevation of -34.2 feet MLLW. At this second location, refusal was encountered at -42.2 feet MLLW due to red/gray stiff clay (native material). Therefore, both a surface (D-ATw-S-2) and a clay (D-ATw-C-2) sample were collected.

D-ATw-S-3/D-ATw-C-4. Similar to previous station, the length of core required to reach the project depth of -44 feet MLLW was longer than could be reached with a 20-foot core barre (targe penetration = 25.2 feet). The "stair-step" approach was required at this station because the one penetration length at the target location was 16.2 feet (bottom core elevation of -35.0 feet MLLW) but did not reach native material. Therefore, EPA was consulted EPA on-station reported to use the "stair-step" approach. ANAMAR were able to find a location downshop of the target location with a top of core elevation of -34.4 feet MLLW. At this second downshop of the target location with a top of core elevation of -34.4 feet MLLW. At this second downshop of the target location with a top of core elevation of -34.4 feet MLLW. At this second downshop of the target location, refusal was encountered at -43 feet MLLW, and there was change in stratification (naive miterial) consisting of sand/clay, large shells, and rocks. Therefore, both a surface (D-ATW-C-4) sample were collected.

D-ATw-S-4/D-ATw-C-3. Refusal was encountered (between 20.4 and -21.4 feet MLLW) above the project depth of -44 feet MLLW due to hard, stiff clay, native clay material was encountered at a much shallower elevation at this station compared to the other three stations in this reach. Two cores were collected from this station to get adequate volume of material for the surface composite sample. Therefore, both a surface (D-ATw-S-4) and a clay (D-ATw-C-3) sample were collected.

Compositing Note: Given that the care lengths of maintenance material varied significantly between the four subsample stations with the Army Terminal Widener Reach, EPA advised the field team to mix proportional volumes based on feet of material recovered from the four subsamples for the composite. Spose columes were calculated and provided to the compositing team.

San Antonio Extension San mary

D-SAx-S-1/D-B-D-7. Second to levation (-40.2 feet) at this station is below the project depth of -38 feet MLLW. EPA was consulted and advised ANAMAR to use a grab sampler to collect unconsolidated...pinterance material at the surface. Therefore, no deepening sample (D-B-D-7) was collected.

D-SAx-S-2/D-B-D-8. Sediment elevation (-38.1 feet) at this station is below project depth of -38 het MLLW. EPA was consulted and advised ANAMAR to use a grab sampler to collect unconsolitated maintenance material at the surface. Therefore, no deepening sample (D-B-D-8) was considered.

D SA 2-3/D-B-D-9. Project depth of -38 feet MLLW was reached at this station. Two cores were collected at this station, and no deepening (native) material was encountered. Therefore, o deepening sample (D-B-D-9) was collected.

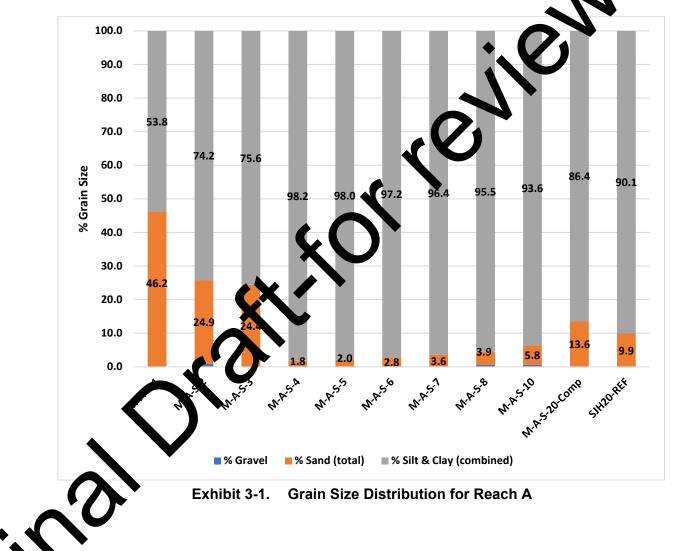


3.2 Physical Testing Data

Grain size distribution and total solids were analyzed in project composite samples, subsamples, individual clay/native material samples, and the reference sample. The following parameters were also analyzed for the composite sample: bulk density, specific gravity, and Atterberg limits. Results are presented in Tables 3 through 5.

San Juan Harbor Maintenance Reach A

Subsamples and the composite sample from Reach A stations were primarily composed rangegrained material (silt/clay) ranging from 53.8% to 98.2%. Exhibit 3-1 shows a bar grap (of th grain size results. The U.S. Soil Classification System (USCS) class was either CH (clay 1 bin plasticity, elastic silt) or MH (silt of high plasticity, elastic silt). Complete results are presented in Tables 3 and 5.





San Juan Harbor Maintenance Reach B

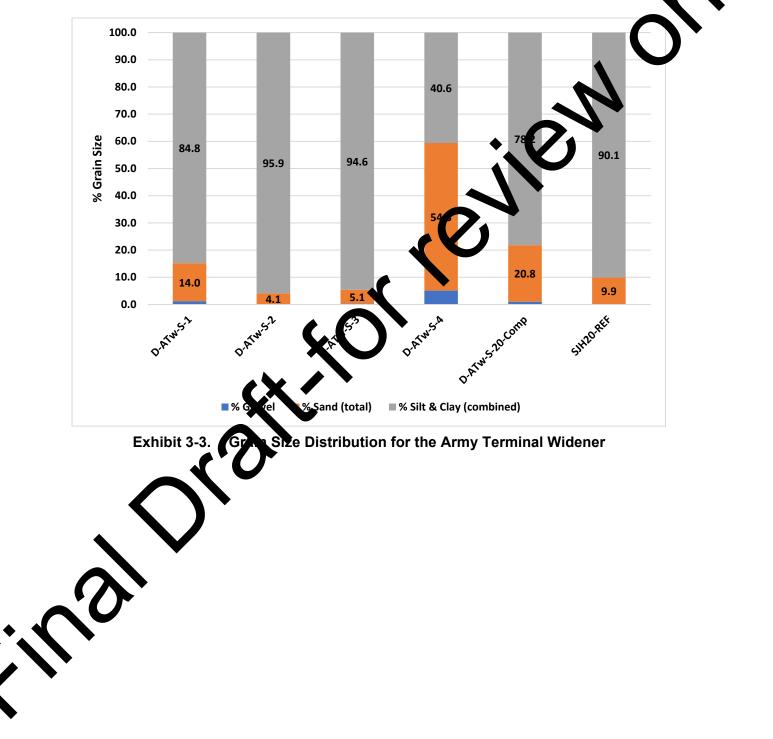
With the exception of subsample M-B-S-2, all subsamples and the composite sample from Reach B stations were primarily composed of fine-grained material (silt/clay) ranging from 52.4% to 95.0%. M-B-S-2 was composed primarily of sand (57.7%). Exhibit 3-2 shows a bar graph of the grain size results. The USCS class was either CH (clay of high plasticity, elastic silt) or SC (clayey sand). Complete results are presented in Tables 3 and 5.





Army Terminal Widener Reach

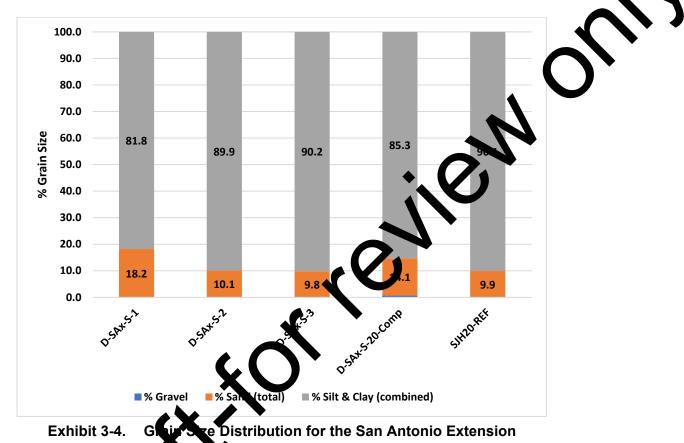
With the exception of subsample D-ATw-S-4, all subsamples and the composite sample from the Army Terminal Widener Reach were primarily composed of fine-grained material (silt/clay) ranging from 78.2% to 95.9%. D-ATw-S-4 was composed primarily of sand (54.3%) with 5.1% gravel. Exhibit 3-3 shows a bar graph of the grain size results. The USCS class was either CH (clay of high plasticity, elastic silt) or SC (clayey sand). Complete results are presented in Tables 3 and 5.





San Antonio Extension

All subsamples and the composite sample from the San Antonio Extension Reach were primarily composed of fine-grained material (silt/clay) ranging from 81.8% to 90.2%. Exhibit 3-4 shows a bar graph of the grain size results. The USCS class was CH (clay of high plasticity, elastic silt). Complete results are presented in Tables 3 and 5.



Individual Clay/Native Material Samples

Samples of stiff clay or meter an epresentative of native/new work material were collected at five locations: one from Reach L and four from the Army Terminal Widener. Exhibit 3-5 shows a bar graph of the grant ize results. Complete results are presented in Table 4.

Sample D-EC-2 (collocated with station M-A-S-3) from the Entrance Channel in Reach A was primarily composed of fine material with 96.7% silt/clay. The USCS class was CH (clay of high plasticity elastic sit).

Sample D-AFw-C-1 (co-located with station D-ATw-S-1) in the Army Terminal Widener was primarily composed of fine material with 83.4% silt/clay. The USCS class was CH (clay of high pustice), elastic silt).

ample D-ATw-C-2 (co-located with station D-ATw-S-2) in the Army Terminal Widener was primarily composed of fine material with 66.0% silt/clay. The USCS class was CH (clay of high plasticity, elastic silt).



Sample D-ATw-C-3 (co-located with station D-ATw-S-4) in the Army Terminal Widener was primarily composed of fine material with 54.7% silt/clay. The USCS class was CH (clay of high plasticity, elastic silt).

Sample D-ATw-C-4 (co-located with station D-ATw-S-3) in the Army Terminal Widener was primarily composed of fine material with 59.5% silt/clay. The USCS class was CH (clay of high plasticity, elastic silt).

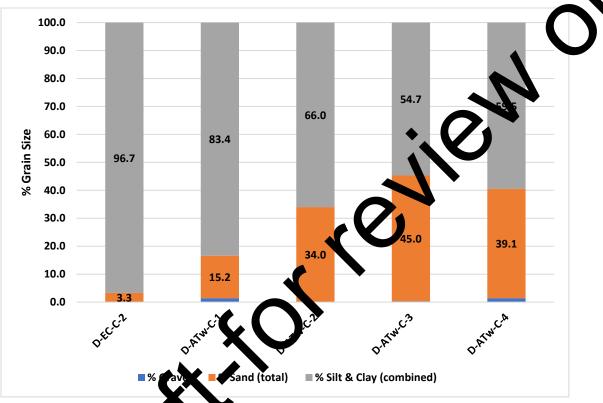


Exhibit 3-5. Grain Size Distribution from Subsamples of Clay/Native Material

3.3 Sediment Clementy

Sediment composite, day/native material samples, and the reference (SJH20-REF) were analyzed for that solids, TOC, metals, pesticides, PAHs, and PCBs. The subsamples were also analyzed for total solids and TOC. Analytical results for sediment chemistry are presented in Tables 1 through 16. Analytical results were compared to the published sediment screening criteria TA and ERL from Buchman (2008). Complete sediment chemistry results are in Aprent. D.

1 otal Solids and TOC

Complete analytical results for total solids and TOC are presented in Tables 6 through 8.

M-A-S-20 composite and subsamples had percent total solids that ranged from 36.90% to 60.23% and TOC concentrations that ranged from 0.84% to 1.96%. Clay/native material sample D-EC-C-2 had percent total solids of 58.43% and a TOC concentration of 1.36%.



M-B-S-20 composite and subsamples had percent total solids that ranged from 48.78% to 61.98% and TOC concentrations that ranged from 0.53% to 1.92%.

D-ATw-S-20 composite and subsamples had percent total solids that ranged from 54.57% to , 65.98% and TOC concentrations that ranged from 0.71% to 1.56%. Clay/native material samples had percent total solids ranging from 65.56% to 78.44%, and TOC concentrations ranging from 0.51% to 0.58%.

D-SAx-S-20 composite and subsamples had percent total solids that ranged from 38, 9% 43.52% and TOC concentrations that ranged from 1.75% to 3.26%.

The reference had 53.83% total solids and 0.90% TOC.

3.3.2 Metals

All nine metals analyzed were detected in concentrations above the NoL A all of the project composite samples. With the exception of cadmium, all other metals analyzed were also detected in concentrations above the MDL in the reference and several of the induitiduarelay/native material subsamples. Exhibit 3-6 summarizes the analytical results for metals in rediment compared to the TEL and ERL. Complete results are provided in Tables 9 and 12

Composite Samples

M-A-S-20-COMP had concentrations of arsenic, copper mercury, and nickel that exceeded the TEL and (or) ERL. M-B-S-20-COMP had concentrations of arsenic, copper, and mercury that exceeded the TEL and (or) ERL. D-ATw-S-20-COMP had concentrations of arsenic, copper, and nickel that exceeded the TEL and (or) ERL. D-SN s-S-20 had concentrations of arsenic, copper, lead, mercury, nickel, silver, and zinc that exceeded the TEL and (or) ERL.

Clay/Native Material Samples

D-EC-C-2 had concentrations of arsenic, concer, mercury, and nickel that exceeded the TEL and and ERL. D-ATw-C-1, C-2 and C-4 had concentrations of arsenic and copper that exceeded the TEL and (or) ERL. D-ATw-C-1 had concentrations of copper that exceeded the TEL.

Reference

SJH20-REF had concentration of arsenic, copper, and nickel that exceeded the TEL and (or) the ERL.



					C	Concentratio	n (mg/kg)					
					Sam	ple ID						
			Composit	e Samples			Clay/	Native Subsa	ples			
Analyte	SJH20 REF	M-A-S-20- COMP	M-B-S-20- COMP	D-ATw-S-20- COMP	D-SAx-S-20- COMP	D-EC-C-2	D-ATw-C-1	P-A 1-0	-ATw-C-3	D-ATw-C-4	TEL	ERL
Arsenic	13.6	15.0	11.9	18.4	13.1	16.5	24.4	13.1	6.54	10.5	7.24	8.2
Cadmium	ND	0.15	0.10	0.05	0.29	ND	0.06	08	ND	ND	0.676	1.2
Chromium	46.4	42.0	34.0	49.6	43.9	49.0	8.5	38.8	33.3	33.8	52.3	81
Copper	63.8	66.0	48.0	47.5	90.5	49.2		37.2	22.3	25.1	18.7	34
Lead	16.3	26.1	23.9	9.70	54.6	9.19	5.75	10.3	5.10	4.23	30.24	46.7
Mercury	0.116	0.342	0.371	0.105	2.28	5 1	0.0712	0.127	0.116	0.0375	0.13	0.15
Nickel	29.3	20.4	14.5	23.6	17.7	22	13.2	15.8	6.79	15.1	15.9	20.9
Silver	0.11	0.62	0.45	0.26	62	0 2	0.04	0.28	0.06	0.06	0.73	1
Zinc	73.9	117	85.6	69.6	155	62.9	45.4	55.7	17.5	36.6	124	150

Exhibit 3-6. Summary of Analytical Results for Metals in Sediment Composites and Clay/Native Mature Subsamples

Bolded values exceed the TEL and/or ERL.

Non-detect (ND) = The analyte was not detected at or abd x = No TEL or ERL published for that parameter.

See Tables 9 and 10 for complete results.



3.3.3 Pesticides

Of the 15 pesticides tested, two [o,p' (2,4')-DDE and p,p' (4,4')-DDE] were detected above the MDL (J-qualified or greater) in one or more samples. For dieldrin, no results were greater than the MDL (U-qualified); but the MDL (0.11 μ g/kg) exceeded the ERL of 0.02 μ g/kg for all samples. However, the MDL for dieldrin was below the EPA Region 2 target detection limit of 1 μ g/kg in Table 13-2 of the SAP/QAPP (Appendix A). Results are summarized below, and complete result are provided in Tables 11 and 12.

Composite Samples

M-A-S-20-COMP and M-B-S-20-COMP had a concentration of p,p' (4,4')-DDE that was greater than the MDL (J-qualified) but did not exceed the ERL or the TEL. D-SAx-S-20-COMP had concentrations of o,p' (2,4')-DDE and p,p' (4,4')-DDE that were greater than the MRL, and p,p' (4,4')-DDE concentrations exceeded the ERL and TEL. In samples M-A-S-20-COMP and D-SAx-20-COMP, the MDLs/MRLs for p,p' (4,4')-DDT and dieldrin were elevated the EPA Region 2 target detection limit of 1 μ g/kg. See Subsection 4.4.2.3 and the COAR (Appendix E) for more information. No other pesticides were detected in concentrations are elevated to MDLs (U-qualified).

Clay/Native Material Samples

None of the results for the subsamples were detected in corservations greater than the MDL; all results were U-qualified. The MDLs and MRLs met the EV/ Region 2 target detection limit of $1 \mu g/kg$.

Reference

None of the results for SJH20-REF were detected in concentrations greater than the MDL; all results were U-qualified. However, the MDL for diadrin (0.11 μ g/kg) exceeded the ERL (0.02 μ g/kg). The MDLs and MRLs met the EPA Regio 2 target detection limit of 1 μ g/kg.

3.3.4 PAHs

All of the 16 PAH analytes to technere detected above the MDL (J-qualified or greater) in one or more composites or subsances. Soveral composite samples had concentrations of PAH analytes that exceeded the uplicable TEL or ERL. The MDLs and MRLs met the EPA Region 2 target detection limit of 1.0 r.g. Results per reach are summarized below and in Exhibit 3-7. Complete results are provided in Tables 13 and 14.

Composite ample

In each of the composite samples, all PAH analytes were detected in concentrations greater than the MDL (J-qualified) or MRL with the exception of acenaphthene in M-B-S-20-COMP. In samples M-A-S-R0-COMP, M-B-S-20-COMP, and D-SAx-20-COMP, acenaphthylene and dibenzo(ab)anthracene concentrations exceeded the TEL. In sample D-SAx-20-COMP, benzo(ab)anthracene and total HMW PAHs concentrations exceeded the TEL.

Clay Valve Material Samples

In samples D-EC-C-2 and D-AT-C-2, all but one of the PAH analytes were detected in concentrations greater than the MDL (J-qualified). In sample D-ATw-C-1, none of PAH analytes were detected in concentrations greater than the MDL (U-qualified). In sample D-ATw-C-3, one PAH analyte was detected in concentrations greater than the MDL (J-qualified). In sample DATw-C-4, nine PAH analytes were detected in concentrations greater than the MDL (J-qualified). In sample (J-qualified). In sample DATw-C-4, nine PAH analytes were detected in concentrations greater than the MDL (J-qualified). In sample (J-qualified). None of the results exceeded the TEL or ERL.



Reference

All PAH analytes that were detected in SJH20-REF were in concentrations greater than the MDL (J-qualified). None of the results exceeded the TEL or ERL.

3.3.5 PCBs

Up to 20 of the 22 PCB congeners tested were detected in concentrations above the MDL in one or more samples. All composites, subsamples, and the reference sample had total EPA Region PCB concentrations that exceeded the applicable TEL or ERL. The MDLs met the EPA Region 2 target detection limit of 1 μ g/kg for all samples. The MRL for PCB-5/8 was elevated above the EPA Region 2 target detection limit of 1 μ g/kg because of the co-eluting of the two congener. Results per reach are summarized below and in Exhibit 3-8. Complete results are provided in Tables 15 and 16.

Composite Samples

In sample M-A-S-20-COMP, 10 of the 22 PCB congeners were detected in concentrations greater than the MDL/MRL. In sample M-B-S-20-COMP, eight of the 22 PCB congeners were detected in concentrations greater than the MDL/MRL. In sample D-ATw-S-20-COMP, and the 22 PCB congeners were detected in concentrations greater than the MDL/MRL. In sample D-ATw-S-20-COMP, 20 of the 22 PCB congeners were detected in concentrations greater than the MDL/MRL. All samples had total EPA Region 2 PCB concentrations that exceeded the TEL and ERL.

Clay/Native Material Samples

In samples D-EC-C-2, D-ATw-C-1, D-ATw-C-3, and D-ATw-C-4, none of the PCB congeners were detected in concentrations greater than the viDL (U-qualified). In sample D-ATw-C-2, four of the 22 PCB congeners were detected in concentrations greater than the MDL/MRL. All samples had total EPA Region 2 PCB concentrations that exceeded the TEL and (or) ERL.

Reference

In sample SJH20-REF, none of the 22 RCL congeners were detected in concentrations greater than the MDL (U-qualified). Thereforence had total EPA Region 2 PCB concentrations that exceeded the TEL.



						Concentratio	on (µg/kg)					
					Sam	ple ID						
				e Samples			Clay	/Native Subsa	nple			
Analyte	SJH20 REF	M-A-S-20- COMP	M-B-S-20- COMP	D-ATw-S- 20-COMP	D-SAx-S- 20-COMP	D-EC-C-2	D-ATw-C-1	D-ATw	DC-3	D-ATw-C-4	TEL	ERL
Acenaphthene	1.61	1.97	ND	0.59	4.47	ND	ND		ND	ND	6.71	16
Acenaphthylene	3.60	7.73	17.8	1.77	22.1	2.55	ND		ND	ND	5.87	44
Anthracene	3.74	8.83	16.6	1.14	31.8	2.01	ND	94	ND	ND	46.9	85.3
Benzo(a)anthracene	15.8	32.0	45.2	3.27	68.0	8.51	NE	4.18	ND	0.89	74.8	261
Benzo(a)pyrene	17.8	45.4	88.7	5.00	125	15.1		8.07	ND	1.51	88.8	430
Benzo(b)fluoranthene	12.2	42.6	91.2	4.50	141	13.1		5.65	ND	1.39	х	х
Benzo(g,h,i)perylene	22.5	41.9	80.5	6.10	111	15.3	ND	10.2	ND	1.78	х	х
Benzo(k)fluoranthene	7.31	21.5	48.6	2.51	71.8	7.12	ND	3.09	ND	ND	х	х
Chrysene	15.6	35.1	53.7	3.78	73.7	10.0	ND	4.80	ND	1.31	108	384
Dibenzo(a,h)anthracene	3.61	11.3	20.2	2.01	28.2	4.17	ND	2.53	ND	ND	6.22	63.4
Fluoranthene	21.5	40.8	67.6	3.32	76.8	14	ND	5.67	0.56	1.25	113	600
Fluorene	2.76	4.33	4.12	0.99	7.2	0.91	ND	1.33	ND	ND	21.2	19
Indeno(1,2,3-cd)pyrene	13.7	35.9	70.5	5.27	95.3	11.1	ND	7.92	ND	1.41	х	х
Naphthalene	4.75	4.16	6.47	1.40	6.5	1.79	ND	1.30	ND	ND	34.6	160
Phenanthrene	15.4	20.3	29.5		30.1	3.10	ND	4.05	ND	0.97	86.7	240
Pyrene	29.1	48.8	70.9	5.66	83.9	11.3	ND	8.41	ND	1.76	153	665
Total LMW PAHs	31.9	47.3	75.1	8.1	103	10.9	5.13	10.9	5.14	5.39	312	552
Total HMW PAHs	159	355	637	1.4	875	105	8.70	60.5	8.79	13.0	655	1700
Total PAHs	191	403		.9.7	977	116	13.8	71.4	13.9	18.3	1684	4022

Exhibit 3-7. Summary of Analytical Results for PAHs in Sediment Composites and Clay/Native Material Subsamples

Bolded values exceed the TEL and/or EPT Non-detect (ND) = The analyte was no detected at on above the MDL.

x = No TEL or ERL published for that the rameter.

See Tables 13 and 14 for complete result





						Concentratio	n (µg/kg)					
					Sam	ple ID						
				e Samples			Clay/	Native Subsa	mp. ¥e.			
Analyte	SJH20- REF	M-A-S-20- COMP	M-B-S-20- COMP	D-ATw-S- 20-COMP	D-SAx-S- 20-COMP	D-EC-C-2	D-ATw-C-1	D-ATw-C-2	L 11-C-3	D-ATw-C4	TEL	ERL
PCB-5/8	ND	ND	ND	ND	1.42	ND	ND		ND	ND	х	х
PCB-18	ND	ND	ND	ND	3.29	ND	ND		ND	ND	х	х
PCB-28	ND	ND	ND	ND	1.96	ND	ND	U U	ND	ND	х	х
PCB-44	ND	ND	ND	ND	1.91	ND	ND	ND	ND	ND	х	х
PCB-49	ND	1.51	1.32	ND	5.68	ND	ND	ND	ND	ND	х	х
PCB-52	ND	1.84	1.66	ND	7.78	ND	N	ND	ND	ND	х	х
PCB-66	ND	ND	ND	ND	2.75	ND	ND	ND	ND	ND	х	х
PCB-87	ND	ND	ND	ND	1.68	ND	ND	ND	ND	ND	х	х
PCB-101	ND	2.41	1.69	ND	7.36	▲ ND	ND	ND	ND	ND	х	х
PCB-105	ND	ND	ND	ND	1.89	ND	ND	ND	ND	ND	х	х
PCB-118	ND	1.25	1.24	ND	5.83	Þ	ND	ND	ND	ND	х	х
PCB-128	ND	ND	ND	ND	1.:	NĎ	ND	ND	ND	ND	х	х
PCB-138	ND	5.13	3.03	1.19	14.4	ND	ND	1.54	ND	ND	х	х
PCB-153	ND	8.10	4.64	1.90	100	ND	ND	2.50	ND	ND	х	х
PCB-170	ND	1.73	ND		3.08	ND	ND	ND	ND	ND	х	х
PCB-180	ND	4.08	1.69	.01	6.68	ND	ND	1.45	ND	ND	х	х
PCB-183	ND	1.34	ND		2.28	ND	ND	ND	ND	ND	х	х
PCB-184	ND	ND	ND	ND	ND	ND	ND	ND	ND	ND	х	х
PCB-187	ND	3.68	1	ND	5.65	ND	ND	1.12	ND	ND	х	х
PCB-195	ND	ND		ND	ND	ND	ND	ND	ND	ND	х	х
PCB-206	ND	N'	ND	ND	1.43	ND	ND	ND	ND	ND	х	х
PCB-209	ND		D	ND	4.75	ND	ND	ND	ND	ND	х	х
Total EPA Region 2 PCBs	22.0	43.1	30.5	22.9	101	22.0	22.0	24.4	21.8	22.0	21.6	22.7

Exhibit 3-8. Summary of Analytical Results for PCBs in Sediment Composites and Clay/Native Material Subsamples

Bolded values exceed the TEL a d/or ERL.

Non-detect (ND) = The analysis at detected at or above the MDL.

x = No TEL or ERL published for that parameter.

See Tables 15 and 16 for complete results.



3.4 Elutriate and Water Chemistry

Analytical results for metals, pesticides, and PCBs in site water (SJH20-SW), reference water (SJH20-REF-SW), and elutriates generated from the four project composites are presented in Tables 17 through 19. Results for elutriate and water samples are compared to the CMC from EPA (2006, 2015). The water and elutriate chemistry laboratory case narrative and data are in Appendix D.

3.4.1 Metals

None of the metals analyzed were detected in concentrations greater than the CMC in an elutriate or water sample. All metals except mercury were detected in concentrations greater than the MDL in all composite elutriate samples. All MDLs were below target reporting imits in the SAP/QAPP and below applicable CMCs. Complete results are in Table 17.

3.4.2 Pesticides

None of the 15 pesticides analyzed were detected in concentrations a love the MDL in any elutriate or reference site water sample. Ten of the 15 pesticides analyzed were detected in concentrations above the MDL in the site water sample. All MDLs were below target reporting limits in the SAP/QAPP and below applicable CMCs. Complete result are in Table 18.

3.4.3 PCBs

None of the 22 PCB congeners analyzed were detected a concentrations above the MDL in any elutriate or site water samples (U-qualified). There are no CMCs for the PCB congeners tested. Total EPA Region 2 PCB concentrations were 0.014 ng/L for all elutriate and site water samples. All MDLs/MRLs were below target reporting invits in the SAP/QAPP. Complete results are in Table 19.

3.5 Benthic Bioassays

The benthic tests were performed with the species *Ampelisca abdita* and *Americamysis bahia*. The complete toxicity testing coort is EcoAnalysts (2021) is provided in Appendix G.

3.5.1 Ampelisca abd a

The 10-day benthic text wit *A abdita* was initiated on December 1, 2020, and was validated by 96% mean surfaced in the control sediment, meeting the acceptability criterion of \geq 90% survival. Mean survival for the project sediment composites ranged from 73% to 91%. Survival in the test sample D-A14-S-20- 00MP was statistically different than that of the reference. Mean percent survival was when 70% of the reference (90%), indicating that the test composite met the LPC for dispesal. Mean survival results are summarized in Exhibit 3-9.

Water coality parameters, ammonia concentrations, and other test conditions are summarized in Tables 3-2 through 3-4 of the toxicity report by EcoAnalysts (2021) in Appendix G. A summary table *A. abdita* survival in each replicate and the raw data bench sheets are provided in Appendix A.1 of the toxicity testing report (Appendix G).



Exhibit 3-9. Summary of Survival Data for the 10-Day Benthic Test with Ampelisca abdita

Sample ID	Mean Survival (% ± SD)	Statistically Significantly Less Than Reference?	Meets LPC Criteria (mean % survival within 20% of Reference?)
Control	96 (± 4.2)		
SJH20-REF (reference)	90 (± 6.1)		
M-A-S-20-COMP	91 (± 7.4)	No	Yes
M-B-S-20-COMP	90 (± 10.0)	No	Yes
D-ATw-S-20-COMP	73 (± 10.4)	Yes	Т Х б
D-SAx-S-20-COMP	88 (± 10.4)	No	Yes

SD = standard deviation

Source: Table 3-1 of EcoAnalysts (2021)

3.5.2 Americamysis bahia

The 10-day benthic test with *A. bahia* was initiated on December 6, 5220, and was validated by 90% survival in the control, meeting the acceptability criterian of 290%. Mean survival within the *A. bahia* benthic test ranged from 89% to 96% in the 45 mediated by and was not statistically different than that of the reference. Mean percent survival was within 10% of the reference (94%), indicating that the test composites met the LPC for disposal. Mean survival results for all samples are summarized in Exhibit 3-10.

Water quality parameters, ammonia concentrations, and other test conditions are summarized in Tables 3-6 through 3-8 of the toxicity know by ErbAnalysts (2021) in Appendix G. A summary table of survival in each replicate and the raw data bench sheets are provided in Appendix A.2 of the toxicity testing report (Appendix G).

Exhibit 3-10. Summary of Servive Data for the 10-Day Benthic Test with Americamysis bahia

Sapple ID	Mean Survival (% ± SD)	Statistically Significantly Less Than Reference?	Meets LPC Criteria (mean % survival within 10% of Reference?)
Control	90 (± 3.5)		
SJH20-REF (removile)	94 (± 5.5)		
M-A-S-2 COMP	89 (± 8.9)	No	Yes
M-D-STO-COMP	93 (± 5.7)	No	Yes
D-ATW-SIZO-COMP	96 (± 4.2)	No	Yes
D SAX-S-20-COMP	95 (± 3.5)	No	Yes

D = standard deviation

ource: Table 3-5 of EcoAnalysts (2021)

3.6 Water Column Bioassays

Water column tests were performed with the mysid crustacean *Americamysis bahia* (opossum shrimp), the atherinoid fish *Menidia beryllina* (inland silverside), and the larval life stage of the



bivalve mollusk *Mytilus galloprovincialis* (Mediterranean mussel). The complete toxicity testing report by EcoAnalysts (2021) is provided in Appendix G.

3.6.1 Americamysis bahia

The 96-hour water column tests with *A. bahia* were initiated on December 7, 2020. The mean survival rate in the control treatment was 94%, meeting the acceptability criterion of ≥90% survival. Mean survival in the site water sample was 98%, indicating that the site water was acceptable in testing. Stray mysids jumped out of the water and desiccated on the side of the test channer (one each in M-A-S-20-COMP 10% Replicate 1, M-B-S-20-COMP 10% Replicate 1, and 0-ATv S-20-COMP 50% Replicate 1). These mysids were removed from statistical analysis and the start count adjusted accordingly.

Mean percent survival in the 100% elutriate concentration was \geq 98% for the settiment composites. The estimated LC₅₀ values were >100%. Statistical comparison of the 100% test uncentrations to the control survival resulted in no significant difference. The mean survivorship data are summarized in Exhibit 3-11.

Water quality measurements, ammonia concentrations, and test conditions are in Tables 3-10 through 3-12 of the toxicity testing report (Appendix G). A summary table of survival in each replicate and the raw data bench sheets are in Appendix 7.3 of the toxicity testing report (Appendix G).

Exhibit 3-11.	Summary of Survival Data for	Water Col	unn Tests l	Using Americamysis
	bahia		•	

Sample ID	Concentration (%)	M an Survival (% ± SD)	Statistically Significantly Less Than Control?	LC 50 (%)
Control		94 (± 4.2)		
SHH20-SW (site water)	X	98 (± 4.5)	No	
M-A-S-20-COMP	100	99 (± 2.2)	No	>100
M-B-S-20-COMP	100	98 (± 2.7)	No	>100
D-ATw-S-20-COMP	100	98 (± 2.7)	No	>100
D-SAx-S-20-COMP	100	98 (± 2.7)	No	>100

SD = standard viation

Source: Table 3-9 Econalysts (2021)

3.6.2 Minidia beryllina

The waves column test with *M. beryllina* was initiated December 7, 2020, and was validated by 91% near survival in the control, meeting the acceptability criterion of \geq 90%. Mean percent survival in the site water sample was 98%, indicating that it was acceptable for testing.

Lean percent survival in the 100% elutriate concentration ranged from 84% to 96%. The sestimated LC_{50} values were >100% for the test composites. Statistical comparison of the test treatments to the control survival resulted in no significant difference. The mean survivorship data for all samples are summarized in Exhibit 3-12.



Water quality parameters, ammonia concentrations, and other test conditions are summarized in Tables 3-14 through 3-16 of the toxicity report by EcoAnalysts (2021) in Appendix G. A summary table of survival in each replicate and the raw data bench sheets are provided in Appendix A.4 of the toxicity testing report (Appendix G).

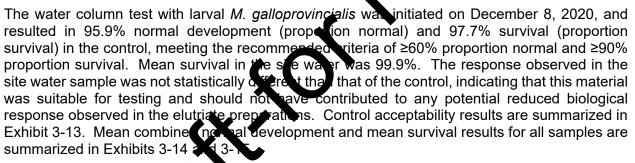
Sample ID	Concentration (%)	Mean Survival (% ± SD)	Statistically Significantly Less Than Control?	
Control		91 (± 11.9)		
SHH20-SW (site water)		98 (± 2.7)	No	N
M-A-S-20-COMP	100	96 (± 4.2)	No	>100
M-B-S-20-COMP	100	92 (± 2.7)	No	>100
D-ATw-S-20-COMP	100	90 (± 6.1)		>100
D-SAx-S-20-COMP	100	84 (± 8.2)	No	>100
2D — standard daviatian				

Exhibit 3-12. Summary of Survival Data for Water Column Tests Using Menidia beryllina

SD = standard deviation

Source: Table 3-13 of EcoAnalysts (2021)

3.6.3 Mytilus galloprovincialis



Water quality parameters and ania concentrations, and other test conditions are summarized in Tables 3-20 through 3.22 or the toxicity report by EcoAnalysts (2021) in Appendix G. A summary table of survival in each represent and the raw data bench sheets are provided in Appendix A.5 of the toxicity tracing report Appendix G).

The estimated C_{50} value for mean proportion normal and proportion survival was >100% for all test settiments, and statistical comparison of the sample results to that of the control resulted in no significant difference.

Exhibit 3 3. *Mytilus galloprovincialis* Control Acceptability Results

Treatment	Mean Proportion Survival (%) ≥90%	Mean Combined Normal Development * ≥60%	Meets Acceptability Criteria?
Control	97.7	95.9	Yes

* Calculated as the total number of normally and abnormally developed embryos ÷ number of embryos stocked (stocking density).

Source: Table 3-17 of EcoAnalysts (2021)



Exhibit 3-14. Mean Combined Normal Development Summary for *Mytilus* galloprovincialis

Sample ID	Concentration (%)	Mean Combined Normal Development * (% ± SD)	Statistically Significantly Less Than Control?	EC ₅₀ (%)	
Control		95.9 (± 4.6)			
SJH20-SW (site water)		94.7 (± 2.8)	No		·
M-A-S-20-COMP	100	98.1 (± 4.2)	No	>100	
M-B-S-20-COMP	100	97.6 (± 4.2)	No	>100	
D-ATw-S-20-COMP	100	99.9 (± 0.2)	No	>100	
D-SAx-S-20-COMP	100	98.0 (± 2.3)	N	>100	

* Calculated as the number of normally developed embryos that survived the duration of indest, number of embryos stocked (stocking density).

SD = standard deviation

Source: Table 3-18 of EcoAnalysts (2021)

Exhibit 3-15. Proportion Survival Summary for Mytilus partoprovincialis

Sample ID	Concentration (%)	Mét Proporti n Survival * V% ± SD)	Statistically Significantly Less Than Control?	LC ₅₀ (%)
Control		97. (± 3.2)		
SJH20-SW (site water)	XV	9.9 (± 0.2)	No	
M-A-S-20-COMP	100	98.6 (± 3.1)	No	>100
M-B-S-20-COMP	10	100.0 (± 0.0)	No	>100
D-ATw-S-20-COMP	X	100.0 (± 0.0)	No	>100
D-SAx-S-20-COMP	100	99.6 (± 0.9)	No	>100

* Calculated as the total number of termally and abnormally developed embryos ÷ number of embryos stocked (stocking density).

SD = standard der an

Source: Table 19 of Eculnal sts (2021)

3.7 Bioaccummation Potential Tests

The 28-cay bioaccumulation tests with *Macoma nasuta* and *Alitta virens* were initiated on December 1a, 2020, respectively. Mean survival in the control was 100% for *M. basuta* and 96.1% for *A. virens*. Mean survival in the reference was 96.8% for *M. nasuta* and 6.0% for *A. virens*. Mean survival in the test composite was \geq 98.4% for *M. nasuta* and \geq 93.0% for *V. virens*. Mean survival results for all samples are summarized in Exhibit 3-16.

Water quality parameters and other test conditions are summarized for the two test species in Tables 3-24 through 3-27 of the toxicity report by EcoAnalysts (2021) in Appendix G. A summary table of survival in each replicate and the raw data bench sheets are provided in Appendices A.6 (for *M. nasuta*) and A.7 (for *A. virens*) of the toxicity testing report (Appendix G).



Exhibit 3-16. Summary of Survival Data for Bioaccumulation Potential Tests Using Macoma nasuta and Alitta virens

	Mean Survival (% ± SD)					
Sample ID	M. nasuta	A. virens				
Control	100 (± 0.0)	96.1 (± 3.5)				
SJH20-REF (reference)	96.8 (± 3.3)	96.0 (± 6.5)				
M-A-S-20-COMP	100 (± 0.0)	99.0 (± 2.2)				
M-B-S-20-COMP	99.2 (± 1.8)	94.0 (± 6.5)				
D-ATw-S-20-COMP	98.4 (± 2.2)	95.0 (± 7_1)				
D-SAx-S-20-COMP	100 (± 0.0)	93.0 <u>(+</u> 7.)				

SD = standard deviation

Source: Table 3-23 of EcoAnalysts (2021)

3.8 Toxicology Summary

Benthic Bioassays



Significant benthic toxicity, relative to the reference treatment, was observed in the *A. abdita* amphipod test for test sample D-ATw-S-20-COMP only. No risnificant toxicity was observed in *A. bahia* mysid test. Mean percent survival in the project composite samples was within the specific test criteria (20% of the reference: amphipod; 10% of the reference: mysid), indicating that the test treatments met the LPC for disposal for the tests.

Water Column Bioassay

No statistically significant toxicity was observed in the 100% elutriate concentrations for the *A. bahia, M. beryllina,* and *M. gallopromicia* is tess.

Bioaccumulation Potential

No significant toxicity was observed in the boaccumulation tests. Survival in the reference and test treatments were ≥93.01, suggesting adequate tissue mass was available for chemical analyses.

3.9 Tissue Chemiltr

Tissue chemistry results for *Masuta* and *A. virens* are presented in Tables 20 through 37. Wet weight tissue memory results for four project samples are compared to the reference (SJH20-REF) and to applicable FDA action levels from FDA (2001, 2011). The laboratory case narrative for tissue chemistry is provided in Appendix D. Complete results of statistical analyses and transformations area. *nasuta* and *A. virens* are provided in Appendix F.

For the weight tables, the laboratory's information management system is not currently able to provide both wet and dry weight concentrations. The results reported were calculated using the wet veight concentration and percent solids provided by the laboratory.

T Lipids and Total Solids in Tissue

Total solids and lipids were analyzed in *M. nasuta* and *A. virens* tissues for the project samples along with the reference and pre-exposure tissues.



<u>Macoma nasuta</u>

Total solids ranged from 16.34% to 18.62% among the project samples, reference, and preexposure tissues. Lipids ranged from 1.5% to 2.5% among these samples. Complete results are in Table 20.

Alitta virens

Total solids ranged from 14.06% to 15.68% among the project samples, reference, and proexposure tissues. Lipids ranged from 2.0% to 3.6% among these samples. Complete results a e in Table 21.

3.9.2 Metals in Tissue

Nine metals were tested in *M. nasuta* and *A. virens* tissues for the project sample, and g with the reference and pre-exposure tissues.

<u>Macoma nasuta</u>

All metals tested were detected in concentrations greater than the MR in the project samples and the reference. Mean concentrations of lead in the project sample M-B-S-20-COMP were statistically significantly greater than those of the reference. Mean concentrations of lead, silver, and zinc in the project sample D-SAx-S-20-COMP were statistically significantly greater than those of the reference. None of the mean concentrations of metals exceeded applicable FDA action levels.

Mean concentrations of metals in *M. nasuta* tissues are summarized in Exhibit 3-17. Complete results are in Tables 22 and 24 for wet weight and the weight metals, respectively. Results of the ToxCalc statistical calculations are provided in Appendix F.

		(X/	Concentrat	ion (mg/kg)		
ļ		ean Sor	centration of F	Replicates	-	
Analyte	M-A-S-20- COMP	В-5 20- Срмр	D-ATw-S- 20-COMP	D-SAx-S- 20-COMP	SJH20-REF (reference)	FDA Action Level
Arsenic	3.31	6.49	3.76	3.89	3.60	86
Cadmium	0.05.4	0.0365	0.0378	0.0380	0.0381	4
Chromium	0.328	0.381	0.340	0.344	0.415	13
Copper		3.70	3.48	3.42	3.41	х
Lead	0.227	0.309	0.154	0.501	0.228	1.7
Mr.(CU)	0.0100	0.0109	0.0095	0.0144	0.0133	1
Nick (0.320	0.370	0.375	0.379	0.450	80
Silver	0.0323	0.0381	0.0375	0.0529	0.0325	х
Zinc	13.2	13.6	14.1	16.0	13.4	х

Exhibit 3-17. Macoma nasuta Tissue: Summary of Mean Wet Weight Metals Results

x = No FDA action level and (or) ecological effects threshold is published for the given parameter.

Bolded values indicate that the mean concentration in project tissues is statistically significantly greater than in the reference tissues, and at least two replicate results are greater than the MDL.

See Table 22 for complete results.



<u>Alitta virens</u>

All metals tested were detected in concentrations greater than the MRL in the project samples and the reference. Mean concentrations of arsenic, cadmium, and chromium in all four project samples were statistically significantly greater than those of the reference. In addition, mean concentrations of copper, nickel, and zinc were statistically significantly greater in D-ATw-S-20-COMP than those of the reference. None of the mean concentrations of metals exceede applicable FDA action levels.

Mean concentrations of metals in *A. virens* tissues are summarized in Exhibit 3-18. Complet results are in Tables 23 and 25 for wet weight and dry weight metals, respectively. Results of the ToxCalc statistical calculations are provided in Appendix F.

			Concentrat	ion (mg/kg)		
		Mean Con	centration of I	Replicates		
Analyte	M-A-S-20- COMP	M-B-S-20- COMP	D-ATw-S- 20-COMP	D-SAx-S- 20-COMP	SJ. 20-REF (reference)	FDA Action Level
Arsenic	2.35	2.53	2.54	200	2.02	76
Cadmium	0.0356	0.0383	0.0355	00206	0.0257	3
Chromium	0.280	0.336	0.325	0.321	0.191	12
Copper	1.31	1.58	2.04	.47	1.47	х
Lead	0.1060	0.135	0.0922	0.130	0.119	1.5
Mercury	0.0188	0.0185	0.021	0.0212	0.0187	1
Nickel	0.135	0.145	160	0.144	0.114	70
Silver	0.0260	0.0193	0089	0.0152	0.0123	Х
Zinc	17.3	(23.)	27.3	20.5	14.5	х

Exhibit 3-18. Alitta virens Tissue: Summary of Mean Wet Weight Metals Res

x = No FDA action level and (or) ecological exects threshold is published for the given parameter.

Bolded values indicate that the more concentration in project tissues is statistically significantly greater than in the reference tissues, and at least two expected results are greater than the MDL. See Table 23 for complete results

3.9.3 Pesticides o Tusue

Fifteen pesticiles were tested in *M. nasuta* and *A. virens* tissues from the four project samples along with the reference and pre-exposure tissues.

Macema hasuta

With the exception of 4,4'-DDE in sample D-SAx-S-20-COMP, none of the pesticides were lated ed in concentrations greater than the MDL in any of the project samples or reference (U-qualified). Mean concentration of 4,4'-DDE (1.49 μ g/kg) in sample D-SAx-S-20-COMP was statistically significantly greater than that of the reference (0.14 μ g/kg). None of the mean concentrations of pesticides exceeded applicable FDA action levels. Complete results are in Tables 26 and 28 for wet weight and dry weight pesticides, respectively. Results of the ToxCalc statistical calculations are provided in Appendix F.



<u>Alitta virens</u>

None of the pesticides were detected in concentrations greater than the MDL in the project samples or the reference. All results were U-qualified. The MDL and MRL for trans-nonachlor were elevated above the target detection limit due to matrix interference. None of the mean concentrations of pesticides exceeded applicable FDA action levels. Complete results are in Tables 27 and 29 for wet weight and dry weight pesticides, respectively. Results of the ToxCal statistical calculations are provided in Appendix F.

3.9.4 PAHs in Tissue

Sixteen PAHs were tested in *M. nasuta* and *A. virens* tissues for the four project samples during with the reference and pre-exposure tissues. Total LMW, total HMW, and total PAHs were calculated from the results of the individual PAHs.

<u>Macoma nasuta</u>

None of the PAHs were detected in concentrations greater than the MDL in the project samples or the reference. All results were U-qualified; therefore, no further statistical analyses or comparisons were needed. Complete results are in Tables 30 and 32 for wet weight and dry weight PAHs, respectively.

Alitta virens

None of the PAHs were detected in concentrations greater than the MDL in the project samples or the reference. All results were U-qualified; therefore, no further statistical analyses or comparisons were needed. Complete results are in Tables 31 and 33 for wet weight and dry weight PAHs, respectively.

3.9.5 PCBs in Tissue

Twenty-two PCB congeners were analyzed in *M. nasuta* and *A. virens* tissues from the four project samples along with the reference and pre-exposure tissues. Total PCBs were calculated from the individual PCB congener equil.

<u>Macoma nasuta</u>

Nine of the PCB conq d were detected above the MRL in at least one of the project ncenterions of PCB congeners 49, 52, 101, 118, 138, and 153 and total sample replicates. EPA Region in some of the project samples were statistically significantly greater than those of the reference Total EPA Region 2 PCB mean concentration in the project samples did ction level. Mean concentrations of PCBs in *M. nasuta* tissues that were not exceed the FDA / statistically sign filly greater than those of the reference are summarized in Exhibit 3-19. Complet results for wet weight and dry weight PCBs are in Tables 34 and 36, respectively. e ToxCalc statistical calculations are provided in Appendix F. Res



Exhibit 3-19. *Macoma nasuta* Tissue: Summary of Mean Wet Weight PCBs Results That Were Statistically Significantly Greater than Those of the Reference

		Concentration (mg/kg)						
		Mean Concentration of Replicates						
Analyte	M-A-S-20- COMP	M-B-S-20- COMP	D-ATw-S- 20-COMP	D-SAx-S- 20-COMP	SJH20-REF (reference)	FDA Action Level		
PCB 49	0.38	0.53	0.40	1.07	0.40			
PCB 52	0.39	0.56	0.40	1.19	0.40			
PCB 101	0.42	0.48	0.40	1.12	0.40	X		
PCB 118	0.38	0.39	0.40	0.68	0.40	х		
PCB 138	0.46	0.44	0.40	0.84	0.40	х		
PCB 153	0.92	0.76	0.40	1.31	0.49	х		
Total EPA Region 2 PCBs	9.11	9.47	8.80	12.6	8.80	2000		

x = No FDA action level and (or) ecological effects threshold is published (at the give parameter.

Bolded values indicate that the mean concentration in project tissues is statistically significantly greater than in the reference tissues, and at least two replicate results are greater than the net. Complete results are in Table 34.

<u>Alitta virens</u>

ove the MRL in at least one of the project Nine of the PCB congeners tested were de sample replicates. Concentrations of CB ers 49, 52, 101, and total EPA Region 2 PCBs bngel in some of the project samples were statically significantly greater than those of the reference. Total EPA Region 2 PCB mean concentrations in the project samples did not exceed the FDA Bs M. nasuta tissues that were statistically significantly action level. Mean concentration of PQ are summarized in Exhibit 3-20. Complete results for wet greater than those of the rel weight and dry weight PCBs bles 35 and 37, respectively. Results of the ToxCalc in statistical calculations are n Appendix F.

Alitta Arens Tigsue: Summary of Mean Wet Weight PCBs Results That
Where Setistically Significantly Greater than Those of the Reference

		····				
			Concentrat	ion (mg/kg)		
		Mean Con	centration of F	Replicates		
Analyte	M-A-S-20- COMP	M-B-S-20- COMP	D-ATw-S- 20-COMP	D-SAx-S- 20-COMP	SJH20-REF (reference)	FDA Action Level
P B / 3	0.40	0.41	0.40	0.54	0.40	х
RCE 52	0.44	0.54	0.40	0.80	0.40	х
PC 101	0.58	0.63	0.40	0.77	0.40	х
Fotal EPA Region 2 PCBs	12.1	12.0	10.6	12.7	11.0	2000

x = No FDA action level and (or) ecological effects threshold is published for the given parameter.

Bolded values indicate that the mean concentration in project tissues is statistically significantly greater than in the reference tissues, and at least two replicate results are greater than the MDL. Complete results are in Table 35.



4 QUALITY ASSURANCE/QUALITY CONTROL

4.1 Coordination with EPA

EPA Region 2 was consulted throughout the sample collection effort for guidance on how to approach sample collection and processing at several stations. Key topics that required consultation involved collection of sample material at stations that required a deepening sample as described in the scope of work and how to collect samples at stations with surface elevations at or below project depth.

General guidelines provided by EPA:

- If shoaling was <2 feet above the target project depth, EPA gave permission to collect the material as a grab sample.
- For the Army Terminal Widener stations, if core length exceeded are longest core barrel available (20 feet), the "stair-step" approach of moving downslop to reach project depth is acceptable, upon final approval by EPA.
- The "clay" or "deepening" samples should represent native (new work) material regardless of the elevation encountered. EPA wanted maintenance (settine) material separated out from the native (new work) material.
- If no native (new work) material was encountered at the project depth, then no "clay" or "deepening" sample was collected at that station
- For Reach B, given that this reach was a nixture of grabs and cores, EPA advised the field team to collect equal volumes from each station for the maintenance (surface) composite sample.
- For Reach B, many stations were blow the deepening project depth. EPA advised the field team to collect a grab sample of anconsolidated material at the surface for the maintenance (surface) composite sample.

A memo was prepared summarizing the field coordination with EPA. A copy of the memo was provided to USACE and EPA and is provided in Appendix J, Pertinent Correspondence.

4.2 Sample Receipt

4.2.1 ARI

Four sediment samples and one site water sample were shipped to ARI on October 27, 2020, and delivered to ARI on October 28, 2020. Sediment and site water for the preparation of elutriates were a livered to MTC on October 29, 2020. All samples were received in good condition and met holding time requirements for both sediment testing and elutriate preparation.

On November 3, 2020, ARI personnel took custody of the reference sediment and site water complex nat were delivered to EcoAnalysts. All samples were received at the laboratories within an avtical holding time and at proper temperature.

2.2 EcoAnalysts

One reference sample, four composite samples, and one site water sample were received in two shipments on October 28 and November 3, 2020. All test samples arrived via two cold boxes at 4.4°C and 4.2°C, respectively, and within the recommended temperature range of 0°C to 6°C upon receipt. Site water and sediment samples were stored in a walk-in cold room at 4 ± 2 °C in



the dark until used for testing. All tests were conducted within the 8-week (56 days) sediment holding time limit.

4.2.3 ALS and Terracon

The cargo container that was used in Puerto Rico was returned to ANAMAR on November 13, 2020. Along with equipment and supplies, the unit contained sediment samples, which ANAMAR packed and shipped to the laboratories on November 16, 2020. Samples shipped to ALS were received on November 18, 2020. Samples were delivered to Terracon on November 19, 2000. All samples were received in good condition.

4.2.4 Tissue Samples

Frozen tissue samples were received at ARI on January 14, 2021, in good conditions Samples were stored in appropriate conditions at the laboratory and thawed to allow pis, at tion only.

4.3 **Physical Analysis**

All physical analyses were performed by Terracon. The analytical result met the quality control criteria specified in the SAP/QAPP.

4.4 Sediment Chemistry

4.4.1 Trace Metals

4.4.1.1 Matrix Spike Recovery

Several spikes were outside control. The laboratory indicates that because the concentration in the sample was substantially higher than in the spike, the accuracy in the spike calculation was reduced.

4.4.1.2 Holding Times

During the initial analysis conducted winin holding time, the recovery for mercury in the standard reference material (SRM) was calculated outside the acceptance range. As a corrective action, samples were frozen and a new SRM was ordered. Re-analysis was performed, and the results were within acceptance chern. It is unlikely that the reported results were substantially affected by the delay in analysis

4.4.2 Pestizion and PCB Congeners

4.4.2.1 Mar ix Spike Recovery

Several pesticities compounds had spike recoveries below 50%, indicating a likely matrix interference. Most results were below the target detection limit and the overall impact on sample results should be low.

Initial and Continuing Calibration Verification

Several compounds had slight exceedances of the acceptance criteria. The overall impact on the sample results was low.

4.4.2.3 Elevated Detection Limits

Pesticide results from samples D-SAx-S-20-COMP and M-A-S-20-COMP had MDLs and MRLs that were above the Region 2 criteria because of matrix interferences. Since the corresponding



tissue samples were analyzed for the affected compounds, the overall impact for these samples was likely to be low.

No other anomalies associated with the analysis of these samples were observed.

4.4.2.4 Standard Reference Material

Endosulfan I for pesticides and several PCB congeners were below the acceptance criteria. Sin the other batch QC were acceptable, the overall impact was likely to be low.

4.4.3 Polycyclic Aromatic Hydrocarbons by EPA Method 8270D

4.4.3.1 Standard Reference Material

All SRM recoveries were within the acceptance limits with the exception of fluore. e, or thracene, and benzo(a)pyrene. Since the remaining batch QC was acceptable, the overall part was likely to be low.

4.4.3.2 Continuing and Initial Calibration Verification

Two verification standards were outside the acceptance criteria; however, the majority were within acceptance limits. Since the exceedances were not significantly outside the acceptance criteria, the overall impact was likely to be low.

4.4.3.3 Spike Recoveries

All spike recoveries were within acceptance criteric with the exception of naphthalene in the spike triplicate. The recovery was consistent with the bike and spike duplicate, indicating a potential matrix interference in the sample.

No other anomalies associated with the manusis of these samples were observed.

4.5 Site Water and Eldviate Chemistry

4.5.1 Trace Metals

4.5.1.1 Matrix Spikes

Cadmium and copper ball solke ecoveries slightly below the acceptance limit, indicating a likely matrix interference.

Note that the spike taget for chromium, lead, and nickel did not meet the criteria specified in the EPA R2 manual. The laboratory indicated that the method could not meet both the low levels needed for reporting limits for at 1 mg/L or lower for copper and silver and the spike target for metals with high reporting limits with minimum levels of 210 mg/L for chromium and 1,050 mg/L for level. The spike recoveries were acceptable for the percent recoveries found.

Festicides and PCB Congeners

1 Matrix Spike Recovery

he matrix spike triplicate was not extracted for SJH20-SW due to a bench sheet error, while D-ATw-S-20-COMP had four matrix spikes samples. All of the samples were batched and had full amounts of batch QC required; however, site-specific QC may be short of spikes.



4.5.2.2 Laboratory Control Standards

One LCS for Endosulfan I was outside the acceptance criteria. All other results were within the acceptance criteria.

No other anomalies associated with the analysis of these samples were observed.

4.6 **Tissue Chemistry**

4.6.1 Trace Metals

4.6.1.1 Matrix Spike Recovery

Spike recoveries were within acceptance limits with the exception of silver for one set of spike triplicates. The relative standard deviation (RSD) for the spike triplicates indicates inwas most likely due to matrix interference in the sample and isolated to silver as all other spike recoveries were acceptable.

4.6.1.2 Continuing Calibration Verification (CCV)

Several CCVs for mercury exceeded the acceptance criteria, but the CCVs analyzed as the next sample were within the acceptance limits, and the samples were bounded by CCVs within the limits.

No other anomalies associated with the analysis of these same were observed.

4.6.2 Pesticides and PCB Congeners

4.6.2.1 Matrix Spike Recovery

Endosulfan sulfate had one spike recovery at 49%, and several spikes for PCB congeners were below the acceptance criteria, indicate a note tial matrix interference in the corresponding samples. All other spikes were acceptable

4.6.2.2 Initial and Continuing Alibration Verification

Several initial calibration verifications (CVs) and CCVs had exceedances for Endosulfan II and PCB 128. The exceedances over an one column only, with the second column having acceptable recoveries; therefore, the sample results were not impacted.

4.6.2.3 Elevater Det ction Limits

Trans-nonablor had elevated detection limits in the *Alitta virens* tissue samples due to matrix interferences. All results were non-detects and the impact is low since all other compounds met the detection limit.

lo my an malies associated with the analysis of these samples were observed.

olycyclic Aromatic Hydrocarbons by EPA Method 8270

Continuing Calibration Verification

everal PAH compounds had exceedances from the acceptance limit but were within the laboratory acceptance criteria. Since all affected sample results were well below the target detection limit, the overall impact to data quality was low.



4.6.3.2 Matrix Spike Recovery

Several compounds had spike recoveries slightly below the acceptance criteria. Since all sample results were well below the target detection limit, the overall impact to data quality was low.

4.6.3.3 Laboratory Control Sample

Several PAH compounds had recoveries below the acceptance criteria, indicating a potential lo f bias in the sample results. The laboratory indicated this was due to multiple cleanup step involved in the preparation of the sample. Since all sample results were well below the target detection limit, the overall impact to data quality was low.

4.7 Toxicology

The quality assurance objectives for toxicity testing are detailed in the Green Loon EPA and USACE 1991) and the laboratory's quality assurance plans. These objectives to accuracy and precision involve all aspects of the testing process, including:

- Water and sediment sampling and handling
- Source and condition of test organisms
- Condition of equipment
- Test conditions
- Instrument calibration
- Use of reference toxicants
- Record-keeping
- Data evaluation

Each test organism was evaluated in renerence toxicant tests during the test period to establish the sensitivity of the test organisms. The reference toxicant LC_{50} or EC_{50} should be within two standard deviations of the distribution laboratory mean. Water quality measurements were monitored to ensure they fell variant rescribed limits.

The methods employed in even phase of the toxicity testing program are detailed in EcoAnalysts' Standard Operating Procedures (SOPs). All EcoAnalysts staff members receive regular, documented training hall SOFs and test methods. All data collected and produced as a result of these analyses were recorded on approved data sheets. If an aspect of a test deviated from protocol, the test was evaluated to determine validity according to the guidance of the regulatory agencies responsible or approval of the proposed permitting action.

4.7.1 Senthic Toxicology Testing

The results of the benthic toxicity tests are presented in this section. The benthic tests were performed with *Ampelisca abdita* and *Americamysis bahia*.

1 Ampelisca abdita

the 10-day benthic test with *A. abdita* was initiated on December 1, 2020, and was validated by 96% survival in the control sample, meeting the acceptability criterion of ≥90%.



Water quality parameters were within acceptable limits throughout the 10-day test, except for pH. While pH was measured at 8.5 in the control treatment, above the targeted range of 7.8 ± 0.5 . It was still within the tolerance range of the test organism and did not negatively affect survival.

The LC₅₀ for the ammonia reference toxicant test was 61.6 mg/L total ammonia and was within two standard deviations of the laboratory mean at the time of testing. This indicates that the test organisms used in this test were of similar sensitivity to those previously tested at the EcoAnalysis laboratory. The concurrent ammonia reference toxicant derived no observed effects concentration (NOEC) values were 34.7 mg/L (total ammonia) and 0.591 mg/L un onized ammonia (UIA). Ammonia concentrations measured within the benthic test were below the ammonia reference toxicant test derived NOEC values for total ammonia and UIA throughout the testing period.

4.7.1.2 Americamysis bahia

The 10-day benthic test with *A. bahia* was initiated on December 8, 2021, and was validated by 90% survival in the control sample, meeting the acceptability criterion of 4.0%

Water quality parameters were within the acceptable limits throughout the 10-day test. Ammonia measurements in overlying water were below the threshold of 0.3 mg² UIA (at pH 7.8) throughout the duration of the test. No afternoon feeding was performed on bay 1 of testing due to a shortage of hatched *Artemia* available for feeding.

The LC₅₀ for the ammonia reference toxicant test was 46.2 mg/L total ammonia and was within two standard deviations of the laboratory mean at the time of testing. This indicates that the test organisms used in this test were of similar sensitivity to those previously tested at the EcoAnalysts laboratory. The concurrent ammonia reference toxicant derived NOEC values were 21.7 mg/L (total ammonia) and 0.380 mg/L (UIA). Alternative derived NOEC values throughout the benthic test were below the ammonia reference to ican test derived NOEC values throughout the testing period.

4.7.2 Water Column Tox tology Testing

The results of the water one on twicity tests are presented in this section. The water column tests were performed with must shripp (*A. bahia*), inland silverside fish (*M. beryllina*), and larvae of the mussel *M. gallebrovicia* is.

4.7.2.1 *Inerical yslo bahia*

The water commuted with *A. bahia* was initiated on December 7, 2020. The mysid test was validated by 94% mean survival in the seawater control, meeting the acceptability criterion of \geq 90%. Itean percent survival in the site water sample was 98%, indicating that the site water was exercised by the site water sample was 98%.

Wate quality parameters were within target limits throughout the duration of the 96-hour test, except for dissolved oxygen. While dissolved oxygen levels fell below the targeted range of >4.0 mg/L on the final day of testing (measured at 3.8 mg/L), the high rate of survival observed in a test treatments indicated that it did not cause any detrimental effects to the test organisms. No eafternoon feeding was performed on Day 2 of testing due to a shortage of hatched *Artemia* available for feeding.



The LC₅₀ for the ammonia reference toxicant test was 46.3 mg/L total ammonia and was within two standard deviations of the laboratory mean at the time of testing. This indicates that the organisms obtained from this supplier were similar in sensitivity to those previously tested at the EcoAnalysts laboratory. The NOEC values were 24.7 mg/L total ammonia and 0.546 mg/L UIA.

4.7.2.2 <u>Menidia beryllina</u>

The water column test with *M. beryllina* was initiated on December 7, 2020. The test was validated by 91% mean survival in the control, meeting the acceptability criterion of \geq 90% durant percent survival in the site water sample was 98%, indicating that the site water was acceptable for testing.

Water quality parameters were within target limits throughout the duration of the the hear test. No feeding was performed on Day 2 of testing due to a shortage of hatched *Artemia*.

The LC₅₀ for the ammonia reference toxicant test was 37.5 mg/L total annunia and was within two standard deviations of the laboratory mean at the time of testing. Basic onchese results, the organisms obtained from this supplier appear to be similar in sensitivity is those previously tested at the EcoAnalysts laboratory. The NOEC values were 29.0 mg/L total ammonia and 0.618 mg/L UIA.

4.7.2.3 <u>Mytilus galloprovincialis</u>

The water column test with *M. galloprovincialis* was interted on December 8, 2020. The larval mussel test resulted in 95.9% normal development (combined proportion normal, number normal ÷ initial number) and 97.7% survival (proportion survival) in the control, meeting the recommended criteria of $\geq 60\%$ proportion proparties and $\geq 90\%$ proportion survival. The embryo stocking density was 24.4 embryos/mt of test solution, within the recommended density of 20 to 30 embryos/mL. Mean survival in the set water was 100%. The response observed in the site water sample was not statistically significantly different than that of the control, indicating that this material was suitable for testing and should not have contributed to any potential reduced biological response observed interval elutriate preparations.

All water quality parameters were within the target limits throughout the duration of the 48-hour test. There was a significant amount of debris in 3 replicates of the 1% concentration and 1 replicate of the 10% concentration of sample D-ATw-S-20-COMP, which was indicative of vial contamination.

The EC₅₀ for the ammunia reference toxicant test was 7.8 mg/L total ammonia and was within two standard deviations of the laboratory mean. This indicates that the population of test organisms used in his test was similar in sensitivity to those previously tested at the EcoAnalysts laboratory. The NOEC values were 5.8 mg/L total ammonia and 0.141 mg/L UIA.

Boaccumulation Tests

The 26-day bioaccumulation tests with *A. virens* and *M. nasuta* were initiated on December 14 and December 9, 2020, respectively. Mean survival in the control samples was 96.1% for *A. virens* and 100% for *M. nasuta*. Reference survival was 96.0% for *A. virens* and 96.8% for *M. nasuta*.

All water quality parameters were within the target limits throughout the duration of the 28-day exposure, except for pH in the *A. virens* test and salinity in the *M. nasuta* test. In the *A. virens*



test, pH was measured below the targeted range, at a minimum of 7.0, in 2 chambers. Water flow was increased in both chambers, and the pH subsequently increased to fall within the target range. Survival remained high in all test treatments. Salinity was measured below the targeted range at 27 ppt during depuration in one chamber of the *M. nasuta* test but was still within the tolerance range of the test organism and would not be expected to influence test results. Inadvertently, only 15 worms rather than 20 were added to Control Replicate 2. As the control tissues are not being analyzed for chemistry, this deviation was not expected to affect the result. The flow rate target per 30 seconds was incorrectly calculated, resulting in flow adjustmente that exceeded the target range of 6 ± 1 volume exchanges per day.

The LC₅₀ for the *A. virens* sodium dodecyl sulfate (SDS) reference toxicant test was 36.8 mg/L SDS and was within two standard deviations of the laboratory mean at the time of testing. The LC₅₀ for the *M. nasuta* reference toxicant test was 39.9 mg/L SDS and was within two standard deviations of the laboratory mean at the time of testing. These reference-toxicant tests indicated that the populations of test organisms used in this study were similar in the study to those previously tested at the EcoAnalysts laboratory.



5 ADDAMS MODEL

Simulations of the STFATE module of the ADDAMS model were run to establish the compliance of the water column toxicity for the San Juan Harbor sediment samples. Each sediment sample represents a separate channel reach or extension. Based on analytical results, no samples were selected for modeling Tier II Water Quality Criteria as all results were below the CMC (National Recommended Water Quality Criteria [EPA 2006, 2015]).

Based on the EC₅₀ results, eight applications (runs) of the models are presented in this report for Section 103 Regulatory Analysis for Ocean Water, Tier III, Short-Term Fate of Dredged Material from Split Hull Barge or Hopper/Toxicity Run.

Results for all the water column toxicology tests show that LC_{50}/EC_{50} 00% across the three species tested for all four San Juan Harbor samples. The project s were modeled to confirm acceptable dilution of the material during disposal to met . STFATE model input parameters used in the module are shown in Exhibits 5-1 t 5-7. The sediment roug physical characteristics (presented in Table 5) for all composite es were used to calculate the volumetric fractions. Values underlined and shown with a haded yellow background were provided by the toxicology laboratory, and the dilution recalculated to allow entry into the simulation (Exhibit 5-7). The files used in the mode runs contained within Appendix H.

Evaluation Type: Tier III, Compare Toxicity Results

Exhibit 5-1. Simulation Type: Descent, Collapse, and Diffusion

Cefficer	its	
Parameter	Keyword	Value
Settling Coefficient	BETA	0.000*
Apparent Mass Coefficient	CM	1.000*
Drag Coefficient	CD	0.500*
Form Drag for Collapsing Crou	CDRAG	1.000*
Skin Friction for Collapsing Couc	CFRIC	0.010*
Drag for an Ellipsoidal Vedge	CD3	0.100*
Drag for a Plate	CD4	1.000*
Friction Between Cloud and Bottom	FRICTN	0.010*
4/3 Low Horizon: Unifusion Dissipation Factor	ALAMDA	0.001*
Unstration d Water Vertical Diffusion Coefficient	AKYO	Pritchard Expression
Closer, mboot Density Gradient Ratio	GAMA	0.250*
Turbuen Chermal Entrainment	ALPHAO	0.235*
Entrationent in Collapse	ALPHAC	0.100*
Stripping Factor	CSTRIP	0.003*

Model default value



Exhibit 5-2. Site Description

Parameter	Value	Units
Number of Grid Points (left to right)	96	n/a
Number of Grid Points (top to bottom)	96	n/a
Spacing Between Grid Points (left to right)	200	ft 🔶
Spacing Between Grid Points (top to bottom)	200	ft
Constant Water Depth	965	ft
Roughness Height at Bottom of Disposal Site	0.005*	ft
Slope of Bottom in X-Direction	0	deg.
Slope of Bottom in Z-Direction	0	deg.
Number of Points in Ambient Density Profile Point	3	n/a
Ambient Density at Depth = 0 ft	1.0236	g/cc
Ambient Density at Depth = 200 ft	1/2-	g/cc
Ambient Density at Depth = 965 ft	♦ 10219	g/cc
Distance from the Top Edge of Grid (upper left corner of site)	6, 90	ft
Distance from the Left Edge of Grid (upper left corner of site)	2,800	ft
Distance from the Top Edge of Grid (lower right corner of site)	12,500	ft
Distance from the Left Edge of Grid (lower right corner of site)	18,800	ft
Number of Depths for Transport-Diffusion Output	<u>3 (0, 450 and 960)</u>	<u>#</u>
		<u></u>

* Model default value

Exhibit 5-3. Current Velocity Data

Р	arameter	Value	Units
X-Direction Velocity		0	ft/sec
Z-Direction Velocity		-1	ft/sec

Exhibit 5-4. Material Data

Parameter	Value	Units
Dredging Site Water Density (a) rage	1.022	g/cc
Number of Layers	1	n/a
Material Velocity at Disposal (X Dir.)	0	ft/s
Material Velocity at Discosal Z-DIX)	-13.5	ft/s

Exhibit 5-5. Output cotions

	Parameter	Value	Units
Duration of Sin Vation		14,400	seconds
Long-term Time Cop		600	seconds

Exhibit 5-6 Disposal Operation Data

	Parameter	Value, Barge/Scow	Unit
	L ηgof Disposal Vessel	200	ft
ヽ ヽ ヽ	Witch of Disposal Vessel	50	ft
	Pre-Disposal Draft	18	ft
	Post-Disposal Draft	5	ft
	Time Needed to Empty the Disposal Bin	20	seconds
	Material Volume	4,800	су
	Location of Disposal from Top of Grid	9,500	ft
	Location of Disposal from Left Edge of Grid	15,800	ft



	M-A-S-20	-COMP	M-B-S-20	-COMP
Analyte	Hopper/Cutter	Mechanical	Hopper/Cutter	Mechanical
Volumetric fractions - Clumps	0.22611	0.60297	0.26789	0.68269
Volumetric fractions - Coarse	0.00273	0.00729	0.00635	0.01
Volumetric fractions - Silt	0.00591	0.01576	0.00476	0.0 212
Volumetric fractions - Clay	0.01146	0.03056	0.00880	0.0224
Solids, %	48.	7	54	6
Specific gravity	2.6	0	2.	¹
Liquid limit	88	1		
<u>LC50/EC50</u>	<u>>100</u>		▲ 10	0
Conc. required to meet criteria	<u>1.00</u>		1.0	0
Dilution required to meet criteria	100)	100	

Exhibit 5-7. Volumetric Fractions and Toxicity Criteria of Dredge Material

	D-ATw-S-2	0-COM	D-SAx-S-20-COMP			
Analyte	Hopper/Cutter	Mechanical	Hopper/Cutter	Mechanical		
Volumetric fractions - Clumps	0.32000	780	0.13836	0.32284		
Volumetric fractions - Coarse	0.00000	0.0000	0.01505	0.03511		
Volumetric fractions - Silt	0.00000	0.00000	0.03715	0.08669		
Volumetric fractions - Clay	0.000	0.00000	0.05015	0.11702		
Solids, %	5.7	7	42.6			
Specific gravity		2	2.61			
Liquid limit	83		92			
LC ₅₀ /EC ₅₀	<u>>10</u>	<u>0</u>	<u>>100</u>			
Conc. required to meet criteria	<u>1.00</u>	<u>)</u>	<u>1.00</u>			
Dilution required to meet criterin	<u>100</u>)	<u>100</u>			

Notes: **Bolded and italicized partmeters** were calculated from Table 5 of this report. <u>Values underlined and shown</u> with a yellow shaded back cound ver provided by the toxicology laboratory, and the dilution required was calculated to allow entry into the simulation. Volumetric fractions were determined using a spreadsheet developed at ERDC. The spreadsheet is *roovided* at the appendices with the filename *SJH volumetric fractions from ERDC calculator.xls*.

Results of the initial mixing simulations after 4 hours of mixing (specified for water column evaluation) and the maximum concentration found outside the disposal area for each dredging unit are summarized in Exhibit 5-8. The location of the maximum concentration is shown as X location and Z location. Input and output files are provided in Appendix H.



Four Hour Disposal Criteria			Disposal Boundary Criteria				
Depth, feet	% Max Conc Above Background on Grid	Dilution on Grid (Da-tox)	X Location	Z Location	Time, hours	Max Conc Outside Disposal Area	Dilutio (Dator)
Sample		A-S-20-COMP					
0	6.70E-40	>10,000	7,200	200	0.50	1.12E-38	> 0,000
450	2.99E-04	>10,000	9,400	1,000	4.0	2.99E-0	>10,000
513 (max)	3.66E-02	2731	9,400	1,000	0.83	3.98E-01	250
960	6.70E-40	>10,000	7,000	200	0.50	1.12 5.	>10,000
Sample	M-A	-S-20-COMP	Hopper Dre	dge (15,000	cubic ya	rds/10. dl.	
0	9.68E-40	>10,000	7,200	200	0.50	.59E-38	>10,000
450	1.05E-14	>10,000	9,400	1,000	4 00	1.03E-14	>10,000
858 (max)	5.29E-02	1889	9,400	1,000	0.8	J.38E-01	185
960	8.59E-03	>10,000	9,400	1,000	0.83	8.83E-02	1132
Sample		S-20-COMP	Clamshell D	redge (4.800	ic ya	ards/load)	
0	1.73E-40	>10,000	7,200	200	0.50	2.84E-39	>10,000
450	1.62E-24	>10,000	9,400	00	4.00	1.62E-24	>10,000
879 (max)	9.44E-03	>10,000	9,400	1,000	0.83	9.30E-02	1074
960	1.60E-03	>10,000	9400	1,000	0.83	1.60E-02	6249
Sample	М-А-	S-20-COMP C	latishell Dr	edge (15,00	0 cubic y	ards/load)	
0	6.94E-40	>10,000	6, 90	200	0.33	7.27E-39	>10,000
450	6.94E-40	> 0,00	600	200	0.33	7.27E-39	>10,000
928 (max)	3.80E-02	31	,400	1,000	0.83	2.52E-01	396
960	8.49E-03	>10,0_0	9,400	1,000	0.83	6.07E-02	1646
Sample		B-S-2 -COMP	Hopper Dre	edge (4,800	cubic yai	ds/load)	
0	5.91E-4	>10,000	7,000	200	0.50	9.33E-39	>10,000
450	5.90E-06	0,000	9,400	1,000	4.0	5.90E-06	>10,000
537 (max)	3.270	3095	9,400	1,000	0.83	3.36E-01	297
960	5.9 E .0	>10,000	9,400	1,000	0.50	9.33E-39	>10,000
Sample	М-В	S-S-20-COMP	Hopper Dre	dge (15,000	cubic ya	rds/load)	
U	. 20E-39	>10,000	7,000	200	0.50	1.51E-38	>10,000
450	6.30E-21	>10,000	9,400	1,000	4.0	6.30E-21	>10,000
874 (m. x)	5.48E-02	1824	9,400	1,000	0.83	5.25E-01	189
960	9.19E-03	>10,000	7,000	200	0.83	8.92E-02	1120
Sample	M-B-	-S-20-COMP	Clamshell D	redge (4,800) cubic y	ards/load)	
	1.40E-40	>10,000	7,200	200	0.50	1.40E-39	>10,000
450	3.43E-29	>10,000	9,400	1,000	4.0	3.43E-29	>10,000
86 (max)	7.65E-03	>10,000	9,400	1,000	0.83	7.33E-02	1364
960	1.32E-03	>10,000	9,400	1,000	0.83	1.29E-02	7752
Sample	М-В-	S-20-COMP C	lamshell Dr	edge (15,00	0 cubic y	ards/load)	
0	5.41E-40	>10,000	6,600	200	0.33	5.59E-39	>10,000
450	5.41E-40	>10,000	6,600	200	0.33	5.59E-39	>10,000
928 (max)	2.96E-02	3377	9,400	1,000	0.83	1.95E-01	512
960	6.68E-03	>10,000	9,400	1,000	0.83	4.74E-02	2109

Exhibit 5-8. Four-Hour Criteria and Disposal Site Boundary Criteria after Initial Mixing





	Four Hour Disposal Criteria			Disposal Boundary Criteria				
Depth, feet	% Max Conc Above Background on Grid	Dilution on Grid (Da-tox)	X Location	Z Location	Time, hours	Max Conc Outside Disposal Area	Dilutio (Dator)	
Sample		Ax-S-20-COM						
0	5.31E-40	>10,000	6,800	200	0.50	7.26E-39	> 0,000	
450	4.83E-36	>10,000	9,400	1,000	4.0	4.83E-36	>10,000	
734 (max)	2.90E-02	3447	9,400	1,000	0.83	2.695-01	371	
960	9.33E-24	>10,000	9,400	1,000	4.0	9.331	>10,000	
Sample		x-S-20-COMP		edge (15,00			•	
0	1.58E-39	>10,000	6,400	200	0.33	.48E-38	>10,000	
450	1.58E-39	>10,000	6,400	200	P 33	1.4 JE-38	>10,000	
934 (max)	8.67E-02	1152	9,400	1,000	0.57	J,40E-01	184	
960	2.12E-02	4716	9,400	1,000	0.67	1.48E-01	675	
Sample		D-SAx-S-20-COMP Clamshell Dredge (4,80, 5 bic yards/load)						
0	3.41E-40	>10,000	6,400	200	0.33	3.35E-39	>10,000	
450	3.41E-40	>10,000	6,400	0	0.33	3.35E-39	>10,000	
938 (max)	1.87E-02	5347	9,400	1,000	0.83	1.19E-01	839	
960	4.96E-03	>10,000	9 400	1,000	0.83	3.64E-02	>10,000	
Sample		-S-20-COMP				yards/load)		
0	8.96E-40	>10,000	6, 90	200	0.33	7.34E-39	>10,000	
450	8.96E-40	> 10,00	200	200	0.33	7.34E-39	>10,000	
939 (max)	4.91E-02	36	,400	1,000	0.67	2.81E-01	355	
960	1.34E-02	746.	9,400	1,000	0.67	9.08E-02	1100	
Sample		W-S-20-COM					> 10,000	
0	5.71E-4	>10,000	7,200	200	0.50	9.10E-39	>10,000	
450 488 (max)	6.03E-03	3204	9,400	1,000	0.83	1.77E-02 3.27E-01	5649 305	
400 (max) 960	3.172-0	>10,000	9,400 7,200	1,000 200	0.83 0.50	9.10E-39	>10,000	
Sample		w-S-20-COMI	÷				~10,000	
Sample	3.84E-40	>10,000	7,200	200	0.50	1.44E-38	>10,000	
450	2.97E-18	>10,000	9,400	1,000	4.0	2.97E-18	>10,000	
868 (m. K)	5.05E-02	1979	9,400 9,400	1,000	0.83	4.95E-01	201	
960	8.36E-03	>10,000	9,400	1,000	0.83	8.28E-02	1207	
Sample		v-S-20-COMP					1201	
0	1.09E-40	>10,000	7,200	200	0.50	1.81E-39	>10,000	
450	2.17E-23	>10,000	9,400	1,000	4.0	2.17E-23	>10,000	
77 (max)	5.95E-03	>10,000	9,400	1,000	0.83	5.91E-02	1691	
960	1.00E-03	>10,000	9,400	1,000	0.83	1.01E-02	9900	
Sample		-S-20-COMP						
0	4.18E-40	>10,000	6,800	200	0.50	4.36E-39	>10,000	
	4.18E-40	>10,000	6,800	200	0.50	4.36E-39	>10,000	
450								
918 (max)	2.28E-02	4385	9,400	1,000	0.83	1.68E-01	594	

Exhibit 5-8. Four-Hour Criteria and Disposal Site Boundary Criteria after Initial Mixing

Dilution $(D_{a-tox}) = (100 - max conc.)/max conc.$



Conclusion

STFATE modeling was performed using two types of dredging equipment, a clamshell dredge combined with a separate barge or scow and a hopper or cutter dredge. Each type of dredging equipment was modeled with a capacity of 4,800 cubic yards per load based on the largest option currently available in Puerto Rico. The model was also performed with a volume of 15,000 cubb yards per load in case a larger dredging vessel or transport equipment becomes available in the future. All model runs met the disposal criteria for both dredging methods and volume Therefore, the material may be disposed without location or volume restrictions, to a maximum volume of 15,000 cubic yards per load within the ODMDS boundaries in accordance with all criteria specified by EPA Region 2 and USACE Jacksonville District.

Exhibits 5-9 and 5-10 show an aerial map of the ODMDS in relation to the non-recoast of San Juan, Puerto Rico, and a computer-generated image showing specific site details, respectively.

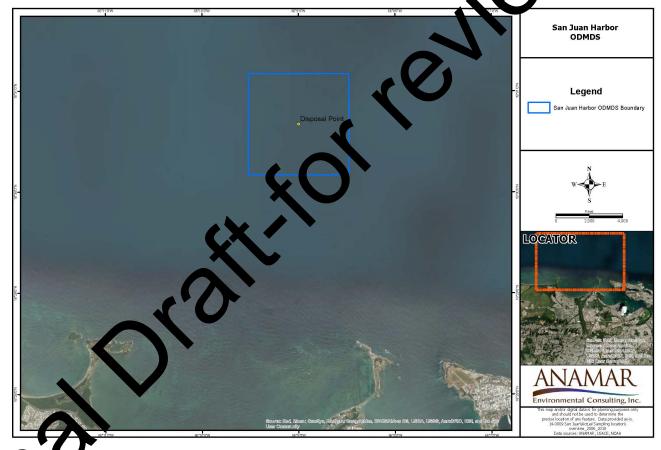


Exhibit 5-9. Aerial Map of San Juan Harbor ODMDS with Disposal Point





es its of the STFATE module of the ADDAMS model indicate that all material from the San Juan bol dredging units may be disposed of at the center of the San Juan Harbor ODMDS using a oper dredge or clamshell with a scow or barge with a carrying capacity of up to 15,000 cubic yards or load without violating applicable disposal criteria.

Exhibit 5-10. Computer-Generated Map of San Juan Harbor ODMDS



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