CCNPP3COLA PEmails

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Cc:	Burkman, Jim; Logan, Carla; Miller, Edward A
Subject:	FW: Proposed e-mail for forwarding UN#10-282 Today
Attachments:	UN#10-282 - Updated Dredging Work Descriptions.pdf
Importance:	High

Attached please find an advance electronic copy of UniStar letter UN#10-282 which was issued today. The advance copy is being forwarded to facilitate discussions in our teleconference schedule for this Friday afternoon, 11/12/10, to discuss NMFS comments and Tidal mitigation.

Regards,

Dimitri Lutchenkov

Director | Environmental Affairs | UniStar Nuclear Energy 410-470-5524 | m 410-370-9090

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Subject:FW: Proposed e-mail for forwarding UN#10-282TodaySent Date:11/11/2010 9:04:26 AMReceived Date:11/11/2010 9:05:25 AMFrom:Lutchenkov, Dimitri			
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Greg Gibson Vice President, Regulatory Affairs 750 East Pratt Street, Suite 1600 Baltimore, Maryland 21202



November 11, 2010

UN#10-282

Jonathan Stewart Tidal Section - Wetlands and Waterways Division Maryland Department of the Environment Water Management Administration 1800 Washington Boulevard Baltimore, Maryland 21230

Woody Francis, Project Manager U.S. Army Corps of Engineers – Baltimore District 10 S. Howard Street Baltimore, Maryland 21201

Subject: Updated Work Description for Tidal Impacts and Dredge Area Sediment Sampling Test Results for Calvert Cliffs Nuclear Power Plant, Unit 3 in Calvert County, Maryland, MDE Project Number 08-WL-1462 (T), 09-NT-0191 (NT), USACE Tracking No. NAB-2007-08123-M05

Enclosure 1 contains updated work descriptions associated with the Joint Federal/State Wetland Permit Application dated 5/16/2008. The updated work descriptions are for tidal impacts only and reflect updated dredge material quantities, a change in pipe cover material for the discharge pipe and fish return, and additional information on the amount of shoreline revetment being impacted by construction.

The dredge material quantity has increased from 58,400 to a total of 66,600 cubic yards which in part reflects the decision not to use dredge material as pipe cover for the fish return and discharge pipe. The amount of material also reflects an increase in dredge material resulting from restoring the barge dock. A breakdown of the dredged material by impact area is as follows:

Intake Area – 1,000 cubic yards

Fish Return – 100 cubic yards

Discharge Pipe – 5,500 cubic yards

Restoration of Barge Unloading Facility - 60,000 cubic yards

Total – 66,600 cubic yards

UN#10-282 Page 2

The area of tidal impact stated in the Joint Federal/State Wetland Permit Application remains the same at 5.7 acres.

Enclosure 2 contains the sediment characterization final report. Physical and chemical analyses of the sediment to be dredged were conducted to evaluate potential placement options.

If you have any questions concerning the attached document, please call Mr. Jim Burkman at (410) 470-5130.

Sincerely,

Greg Gibson

- Enclosures 1) Updated Work Descriptions for Tidal Impact, October 2010
 - 2) Calvert Cliffs Nuclear Power Plant, Unit 3 Project, Sediment Characterization Final Report, EA Engineering, Science, and Technology, Inc., October 2010
- cc: Jim Butch EPA Susan Gray – Power Plant Research Program Cheryl Kerr – MDE John Nichols - NMFS Laura Quinn – NRC Bob Zepp – USFWS Greg Golden - MDNR Kathy Anderson - USACE Bob Sadzinski - MDNR Roland Limpert - MDNR Michael Mansolino - EPA Harriet Nash - NRC Mary Ann Parkhurst - PNL Roy Kropp - PNL

Enclosure 1

Updated Work Descriptions for Tidal Impact October 2010 UN#10-282 Enclosure 1 Page 2 of 3

Work in Tidal Areas:

Work in tidal areas will be restricted to four major areas. The following paragraphs will describe the scope of work in each of these areas.

1. Unit 3 Intake Area

The work in the Unit 3 Intake area will include two major components: a) installation of a permanent sheet pile wall and b) installation of temporary sheet pile wall. The details of the work are shown on Figure 3.

a) Permanent Sheet Pile Wall: A new sheet pile wall will be installed to construct Unit 3 intake area. The sheet pile wall will extend approximately 180 feet from the existing shoreline to the existing baffle wall. The sheet pile wall will be located approximately 90 feet channelward of the approximate mean high water shoreline. The new sheet pile wall along with the existing baffle wall and the shoreline will create a 9,000 square foot wedge-shaped pool. To facilitate installation of the permanent sheet pile wall, approximately 50 feet of the existing shoreline armor protection will be removed. A new armor protection, approximately 75 feet long, will be installed adjacent to the new sheet pile wall. The armor protection, consisting of imported coarse sand / stone, will extend approximately 95 feet channelward.

b) Temporary Sheet Pile Wall: To facilitate the installation of the intake pipes, a temporary sheet pile wall will be installed after removal of the existing (approximately 60 feet along the shoreline) armor protection. The temporary sheet pile wall will extend approximately 35 feet (average) into the wedge-shaped pool. The area within the sheet piling will be dewatered and mechanically dredged to create an approximately 30-feet wide by 35-feet long by 25-feet deep area. Subsequently, extending approximately 20 feet channelward, two 60-inch diameter intake pipes (with trash rack and associated structures) will be installed with the bottom elevation at -25 feet mean low water. Shoreline armor protection (about 10 feet channelward) will be restored as required. The temporary sheet pile wall will then be removed to flood and submerge the intake pipes. The sand and gravel removed by dredging (approximately 1000 cubic yards) will be deposited onsite at an existing upland (non-wetland) environmentally controlled Lake Davies laydown area.

2. Unit 3 Fish Return

A fish return system similar to the existing Unit 1/Unit 2 fish return will be installed as part of the intake structures. The work related to the fish return system in the tidal areas includes installation of a discharge pipe and a rip-rap apron along with associated dredging.

The discharge pipe will consist of an 18-inch diameter high density polyethylene (HDPE) pipe which will be installed approximately 4 feet below the bay bottom in a mechanically excavated trench. The pipe will outfall about 40 feet channelwards from the existing shoreline. The location of the outfall will be protected with a riprap apron. To facilitate installation of the pipe, existing shoreline revetment (approximately 65 feet along the shoreline) will be removed and replaced. This dredging will temporarily impact approximately 100 cubic yards of material. The pipe trench will be filled with imported coarse sand/stone fill. The existing shoreline revetment will be restored to its original configuration after installation of the pipe. Turbidity curtains will be utilized during construction to contain suspended sediments. The sand and gravel removed by dredging will be deposited onsite at an existing upland (non-wetland) environmentally controlled Lake

UN#10-282 Enclosure 1 Page 3 of 3

Davies laydown area.

3. Discharge Outfall Pipe

A 30-inch diameter HDPE pipe with three single port diffusers will be installed in a mechanically excavated trench. The discharge point at the diffusers will be elevated approximately 3 feet above the bay bottom. The pipe will extend approximately 550 feet channelward and will be buried approximately 4 feet below the bay bottom. This burial depth will prevent damage of the pipe from storms and small boat anchors. To facilitate installation of the pipe, existing shoreline revetment (approximately 70 feet along the shoreline) will be removed and replaced. This installation will temporarily impact approximately 38,500 square feet (approximately 0.9 acres) area at the bay bottom. Additionally, a riprap scour pad will be installed at the diffuser outfall permanently impacting 800 square feet area at the bay bottom. The excavated trench will be filled with approximately 5,500 cubic yards of imported coarse sand/stone material. The dredged material (5,500 cubic yards) will be deposited onsite at an existing upland (non-wetland) environmentally controlled Lake Davies laydown area. Turbidity curtains will be utilized during construction to contain suspended sediments.

4. Restoration of Barge Unloading Facility including Maintenance and New Dredging

The existing barge unloading facility is intended to be utilized to receive equipment and materials for the construction of the Unit 3. The existing barge slip will be restored and extended to re-establish use of an approximately 1,500 feet long and 130 feet wide (average) channel. The channel area (about 195,000 square feet) will be dredged to a bottom elevation of -16 feet mean low water. The initial 1,065 feet length of the dredging is considered maintenance dredging. The remaining 435 feet is considered an extension beyond the original dredging limits to reach the bottom elevation of -16 feet mean low water. Of 60,000 cubic yards of total estimated dredging, 57,000 cubic yards are considered maintenance dredging and 3,000 cubic yards are considered maintenance dredging and 3,000 cubic yards are considered in the channel will also be removed.

The dredged material will be deposited at existing onsite upland (non-wetland) environmentally controlled Lake Davies laydown area. Turbidity curtains will be utilized during construction to contain the suspended sediments.

The scope of work in this area will also include maintenance dredging near the shoreline to remove sediments which have mounded up over the past 30 years and restoration of an existing culvert outfall. Due to silt build up over the years, the discharge from this outfall meanders in a north-south direction prior to discharging into the barge slip area. The restoration activities in this area will include the installation of a riprap apron (Figure 6J) in front the existing outfall allowing the discharge to flow directly in the bay as originally designed. The riprap apron will extend approximately 40 feet channelward.

Additionally, a new sheet pile wall will be installed along the shore line in front of the existing bulk head which was built as part of the original design. On the land side of the new sheet pile, a concrete apron will be placed along with a gravel apron to allow equipment to be off-loaded from barges with wheel mounted transporters.

Enclosure 2

Calvert Cliffs Nuclear Power Plant Unit 3 Project Sediment Characterization Final Report EA Engineering, Science, and Technology, Inc. October 2010

CALVERT CLIFFS NUCLEAR POWER PLANT UNIT 3 PROJECT

SEDIMENT CHARACTERIZATION

FINAL REPORT

Prepared for

UniStar Nuclear Energy Calvert Cliffs Nuclear Power Plant 1650 Calvert Cliffs Parkway Lusby, Maryland 20657

Prepared by

EA Engineering, Science, and Technology, Inc. 15 Loveton Circle Sparks, Maryland 21152

October 2010

TABLE OF CONTENTS

1.	PRO.	JECT DESCRIPTION	. 1
	1.1	Project Location	. 1
	1.2	Project Background and Scope	. 1
	1.3	Technical Approach	. 2
2.	SAI	MPLING PROCEDURES	. 5
	2.1	Sampling Locations	. 5
	2.2	In Situ Water Quality Measurements	. 5
	2.3	Site Water and Elutriate Water	. 5
	2.4	Sediment Sampling	. 6
	2.5	Sample Processing and Compositing	. 6
	2.6	Equipment Decontamination Procedures	. 7
	2.7	Equipment Blank	. 8
	2.8	Documentation	. 8
	2.8.1	Numbering System	. 8
	2.8.2	Sample Documentation	. 9
	2.8.3	Chain-of-Custody Records	9
	2.0.5	Cham-of-Custody Records	• /
3.		ALYTICAL METHODS AND DATA EVALUATION	
3.			10
3.	AN	ALYTICAL METHODS AND DATA EVALUATION	10 10
3.	AN 3.1	ALYTICAL METHODS AND DATA EVALUATION Analytical Methods	10 10 11
3.	AN 3.1 3.2	ALYTICAL METHODS AND DATA EVALUATION Analytical Methods Detection Limits	10 10 11 12
3.	AN 3.1 3.2 3.3	ALYTICAL METHODS AND DATA EVALUATION Analytical Methods Detection Limits Laboratory Quality Control Samples	10 10 11 12 12
3.	AN 3.1 3.2 3.3 3.4	ALYTICAL METHODS AND DATA EVALUATION Analytical Methods Detection Limits Laboratory Quality Control Samples Data Analysis	10 10 11 12 12 12
3.	AN 3.1 3.2 3.3 3.4 3.4.1	ALYTICAL METHODS AND DATA EVALUATION Analytical Methods Detection Limits Laboratory Quality Control Samples Data Analysis Calculations for Total PCBs and Total PAHs	 10 10 11 12 12 12 13
3.	AN 3.1 3.2 3.3 3.4 3.4.1 3.4.2	ALYTICAL METHODS AND DATA EVALUATION Analytical Methods Detection Limits Laboratory Quality Control Samples Data Analysis Calculations for Total PCBs and Total PAHs Comparison to Sediment Quality Guidelines	 10 10 11 12 12 12 13 14
3.	AN 3.1 3.2 3.3 3.4 3.4.1 3.4.2 3.4.3	ALYTICAL METHODS AND DATA EVALUATION	 10 10 11 12 12 12 13 14 14
3.	AN 3.1 3.2 3.3 3.4 3.4.1 3.4.2 3.4.3 3.4.4	ALYTICAL METHODS AND DATA EVALUATION	 10 10 11 12 12 12 13 14 14 14
	AN 3.1 3.2 3.3 3.4 3.4.1 3.4.2 3.4.3 3.4.3 3.4.4 3.4.5 3.4.6	ALYTICAL METHODS AND DATA EVALUATION	 10 10 11 12 12 12 13 14 14 14 14
	AN 3.1 3.2 3.3 3.4 3.4.1 3.4.2 3.4.3 3.4.3 3.4.4 3.4.5 3.4.6	ALYTICAL METHODS AND DATA EVALUATION Analytical Methods Detection Limits Laboratory Quality Control Samples Data Analysis Calculations for Total PCBs and Total PAHs Comparison to Sediment Quality Guidelines Comparison to USEPA/State of Maryland Water Quality Criteria Calculation of Acute and Chronic Ammonia (NH ₃ -N) Criteria Evaluation of TCLP Data Comparison to Shirley Plantation Screening Criteria	 10 10 11 12 12 12 13 14 14 14 14 14 15
	AN 3.1 3.2 3.3 3.4 3.4.1 3.4.2 3.4.3 3.4.4 3.4.5 3.4.6 ANA	ALYTICAL METHODS AND DATA EVALUATION Analytical Methods Detection Limits Laboratory Quality Control Samples Data Analysis Calculations for Total PCBs and Total PAHs Comparison to Sediment Quality Guidelines Comparison to USEPA/State of Maryland Water Quality Criteria Calculation of Acute and Chronic Ammonia (NH ₃ -N) Criteria Evaluation of TCLP Data Comparison to Shirley Plantation Screening Criteria LYTICAL RESULTS	 10 11 12 12 12 13 14 14 14 14 15 15
	AN 3.1 3.2 3.3 3.4 3.4.1 3.4.2 3.4.3 3.4.4 3.4.5 3.4.6 ANA 4.1	ALYTICAL METHODS AND DATA EVALUATION Analytical Methods Detection Limits Laboratory Quality Control Samples Data Analysis Calculations for Total PCBs and Total PAHs Comparison to Sediment Quality Guidelines Comparison to USEPA/State of Maryland Water Quality Criteria Calculation of Acute and Chronic Ammonia (NH ₃ -N) Criteria Evaluation of TCLP Data Comparison to Shirley Plantation Screening Criteria LYTICAL RESULTS Bulk Sediment Results	 10 11 12 12 12 13 14 14 14 14 15 15

4.1.4	Organic Constituents	16
4.2	Site Water and Elutriate Results	
4.2.1	General Chemistry Analyses	
4.2.2	Inorganic Constituents	
4.2.3	Organic Constituents	17
4.3	TCLP Results	
4.4	Shirley Plantation Screening	
5. SU	MMARY AND CONCLUSIONS	
5.1	Physical Analyses	
5.2	Chemical Analyses	
5.3	Oyster Survey	
5.4	Conclusions	
6. RE	FERENCES	

LIST OF APPENDICES

Appendix A	Oyster Survey Report
Appendix B	Field Notes
Appendix C	Chain-of-Custody Forms
Appendix D	Project Limits

LIST OF FIGURES

Figure 1. Project Location Map

Figure 2. Proposed Dredging Areas and Project Sampling Locations

LIST OF TABLES

- Table 1.Sampling Locations and Coring Summary
- Table 2.In Situ Water Quality Data
- Table 3.Required Containers, Preservation Techniques, and Holding Times for Aqueous
Samples
- Table 4.Required Containers, Preservation Techniques, and Holding Times for Sediment
Samples
- Table 5.Analytical Methods
- Table 6.Marine Sediment Benchmarks
- Table 7.USEPA and State of Maryland Acute and Chronic Saltwater Quality Criteria for
Aquatic Life
- Table 8.TCLP Regulatory Guidelines
- Table 9.Shirley Plantation Screening Table
- Table 10.Physical Characteristics of Sediment
- Table 11.General Chemistry Concentrations in Sediment
- Table 12.Metal Concentrations (mg/kg) in Sediment
- Table 13.PAH Concentrations (µg/kg) in Sediment
- Table 14.PCB Congener Concentrations (µg/kg) in Sediment
- Table 15.PCB Aroclor Concentrations (µg/kg) in Sediment
- Table 16.Chlorinated Pesticide Concentrations (µg/kg) in Sediment
- Table 17. SVOC Concentrations (μ g/kg) in Sediment
- Table 18. Butyltin and 2,3,7,8-TCDD Concentrations (µg/kg) in Sediment
- Table 19.General Chemistry Concentrations in Site Water and Standard and Effluent
Elutriates
- Table 20. Metal Concentrations (µg/L) in Site Water and Standard and Effluent Elutriates
- Table 21. PAH Concentrations (µg/L) in Site Water and Standard and Effluent Elutriates
- Table 22.PCB Congener Concentrations ($\mu g/L$) in Site Water and Standard and Effluent
Elutriates
- Table 23.PCB Aroclor Concentrations ($\mu g/L$) in Site Water and Standard and Effluent
Elutriates
- Table 24.Chlorinated Pesticide Concentrations (μ g/L) in Site Water and Standard and
Effluent Elutriates
- Table 25. SVOC Concentrations (µg/L) in Site Water and Standard and Effluent Elutriates
- Table 26.Butyltin and 2,3,7,8-TCDD Concentrations (μ g/L) in Site Water and Standard and
Effluent Elutriates
- Table 27.TCLP in Sediment

1. PROJECT DESCRIPTION

EA Engineering, Science, and Technology, Inc. (EA) was contracted by UniStar Nuclear Energy (UniStar) to collect sediment samples offshore of Calvert Cliffs Nuclear Power Plant (CCNPP) in Calvert County, Maryland (Figure 1). Physical and chemical analyses of the sediment to be dredged were conducted to evaluate potential placement options.

The proposed dredging project is part of a UniStar's proposal to develop a third unit at CCNPP. UniStar would be bringing supplies and equipment to the facility via an existing barge unloading facility. Dredging is needed to restore the existing barge dock. UniStar also needs to dredge small areas associated with other infrastructure upgrades and installations. These include installation of a discharge pipe and installation of a fish return. UniStar estimates that 66,000 cubic yards of material will be dredged from the following four areas (Figure 2):

- Barge Unloading Facility
- Wedge Shaped Pool (Intake Area)
- Discharge Pipe
- Fish Return

The objective of the sampling effort was to obtain and analyze sediment and water samples representative of material proposed for dredging as part of the proposed dredging areas. The resulting geotechnical and analytical data were used to characterize the sediments and to determine the suitability of different types of placement options.

1.1 **Project Location**

CCNPP is located in the mid-Chesapeake Bay, north of the confluence of the Patuxent River with the Chesapeake Bay. Dredging activities will occur adjacent to the shoreline of the CCNPP site, which is in Calvert County, Maryland, near Lusby, Maryland.

1.2 Project Background and Scope

UniStar, on behalf of Calvert Cliffs 3 Nuclear Project, LLC, has applied for a license from the Nuclear Regulatory Commission (NRC) to construct a third unit at the existing CCNPP. Unit 3 would be constructed adjacent to Units 1 and 2. Unit 3 is proposed to help meet long-term energy demands within the region and would provide 1,600 megawatts of additional power to the grid. UniStar is proposing to use an existing barge unloading facility to bring equipment and supplies to support construction of the proposed Unit 3. Use of the existing barge unloading facility and pier, requires dredging the access channel to a depth of -16 feet MLW to allow sufficient under-keel clearance to accommodate anticipated barge traffic. To support operation of Unit 3, UniStar plans to install a new discharge pipe and a new fish return, both of which would also require dredging during installation. A fourth area, the wedge shaped pool (intake area), would be dredged to support installation of new intake piping.

The removal of approximately 66,000 cubic yards of material over 5.7 acres of Chesapeake Bay bottom is proposed. Sediment quality analyses were conducted to assess the suitability of the material for various placement options. UniStar has identified the upland placement facility at Port Tobacco at Weanack, better known as Shirley Plantation as one of the potential placement options and requested testing to determine the suitability of the material for placement at the site. Other placement options, such as wetland creation and confined disposal, may be evaluated based on the results of the sediment quality analyses.

The testing protocols used for the sediment quality investigation were determined using the general screening requirements for beneficial use, innovative reuse, and upland placement, as well as the specific requirements for placement Shirley Plantation (Section 1.3). The list of target detection limits, sampling methodologies, and sample holding times for the sediment samples were derived from *QA/QC Guidance for Sampling and Analysis of Sediments, Water, and Tissues for Dredged Material Evaluations* (USEPA/USACE 1995). Based on the data collected, the suitability of the material for various placement options was evaluated.

1.3 Technical Approach

This investigation was designed to identify, analyze, and evaluate the physical and chemical characteristics of sediment, site water, and elutriate samples that are representative of the areas proposed for dredging. The sampling and analytical components for this evaluation were derived from the following guidance documents:

- USEPA/USACE, 1998 (EPA-823-B-98-004). Evaluation of Dredged Material Proposed for Discharge in Waters of the U.S>-Testing Manual [Inland Testing Manual (ITM)]
- USEPA/USACE, 1995 (EPA-823-B-95-001). *QA/QC Guidance for Sampling and Analysis of Sediments, Water, and Tissues for Dredged Material Evaluations.*
- Toxicity Characteristic Leaching Procedure (TCLP) Regulatory Guidelines 40 CFR 261.24

A separate investigation was designed and implemented to determine the quantity of oyster shell within the sediment in the proposed dredging footprint. This study was completed by Dr. Kennedy Paynter at the University of Maryland and has been appended to this document as Appendix A.

1.3.1 Field Sampling Program

The field sampling and sample processing program included the following:

- Collecting sediment samples to project depth using a virbracorer from a total of nine locations in project dredging areas (Figure 2)
- Collecting samples from specified locations with positioning accuracy appropriate for project objectives

- Homogenizing sediment samples from multiple locations into composite samples that were submitted for bulk sediment, standard elutriate, and effluent elutriate preparation
- Collecting a site water sample using a peristaltic pump
- Collecting and transferring sediment and site water to appropriate, laboratory-prepared containers and preserving/holding samples for analysis according to protocols that ensure sample integrity
- Collecting and recording *in Situ* water quality (temperature, pH, dissolved oxygen, and salinity) for each sampling location at the time of sample collection

The sample compositing for each channel is summarized below and in Table 1:

- For the barge dock area, a total of four locations were sampled. Two composite samples (CCU3-BAR-SED-1/2 and CCU3-BAR-SED-3/4), each consisting of sediment from two locations, were created and used for the physical and chemical analyses.
- For the discharge pipe area, one location was sampled. The sample (CCU3-DIS-SED) was homogenized and used for physical and chemical analyses.
- One sample was planned for the wedge shaped pool, but was unable to be sampled because of riprap on the sediment surface.

Field sampling is described further in Section 2. Note that the fish return area was not sampled because at the time of sampling, this area was going to be returned to the excavation area after installation of the fish return. This material will now be placed with the rest of the project dredged material.

The oyster shell survey program (Appendix A) included the following:

- Characterizing bottom type within the dredging area
- Determining shell coverage within the dredging area
- Counting live and dead oysters within the dredging area

1.3.2 Analytical Testing

Analytical testing of the bulk sediments, site water, standard elutriates, and effluent elutriates was conducted by TestAmerica located in Pittsburgh, Pennsylvania. Standard elutriates simulate the release of metals and organic constituents in the water column during open water/ocean placement of material. Effluent elutriates simulate the quality of effluent discharged from confined dredged material disposal area during dredging operations. Methodology for the standard and effluent elutriate preparation is provided in the *Inland Testing Manual* (USEPA/USACE 1998). The analytical program included the following tasks:

- Physical analyses (grain size and percent solids) of sediment from two composite sediment samples, created from four channel locations, and one individual sediment sample;
- Preparation of three standard elutriates from two composite sediment samples and one individual sediment sample;
- Preparation of three effluent elutriates from two composite sediment samples and one individual sediment sample;
- Chemical analysis of bulk sediment for the following project-specific target analytes: ammonia, nitrate-nitrite, total cyanide, total Kjeldahl nitrogen (TKN), total organic carbon (TOC), total phosphorus, total sulfide, total petroleum hydrocarbon (TPH) – diesel range organics (DRO), TPH – gasoline range organics (GRO), metals, polyaromatic hydrocarbons (PAHs), polychlorinated biphenyl (PCB) aroclors, PCB congeners, chlorinated pesticides, semivolatile organic compounds (SVOCs), butyltins, and 2,3,7,8-TCDD;
- Chemical analysis of site water, standard elutriates, and effluent elutriates for the following project-specific target analytes: ammonia as nitrogen, dissolved cyanide, dissolved organic carbon, total Kjeldahl nitrogen, TPH DRO, TPH GRO, metals, PAHs, PCB congeners, PCB aroclors, chlorinated pesticides, SVOCs, 2,3,7,8-TCDD, butyltins (site water only), total nitrate/nitrite (site water and standard elutriates only), total sulfide (site water and standard elutriates only), phosphorous as orthophosphate (site water and standard elutriates only), dissolved nitrate/nitrite (site water and effluent elutriates only), and total phosphorus (site water and effluent elutriates only).

In addition to sediment, water, and elutriate samples, quality control (QC) samples were submitted to the laboratory. Matrix spike/matrix spike duplicate (MS/MSD) samples were analyzed. Analytical methods, target analytes, holding times, reporting limits, and laboratory quality assurance/quality control (QA/QC) protocols are described in Chapter 3 of this report.

1.3.3. Data Analysis

Data analysis included the following tasks:

- Chemical concentrations in sediment samples were compared to Sediment Quality Guidelines (SQGs) (Long et al. 1995) and the TCLP screening criteria.
- Chemical concentrations in sediment samples were compared to Shirley Plantation screening criteria.
- Chemical concentrations in site water, standard elutriates, and effluent elutriate samples were compared to U.S. Environmental Protection Agency (USEPA) and the State of Maryland saltwater acute and chronic water quality criteria (WQC) for aquatic life.

Analysis of oyster data was completed by Dr. Paynter at the University of Maryland and his report is included as Appendix A.

2. SAMPLING PROCEDURES

Field activities consisted of sediment and water sampling offshore of CCNPP within the proposed dredging areas. Sampling was conducted at four locations adjacent to the barge dock and one location within the proposed discharge pipe area (Figure 2). An additional sample location was proposed within the wedge shaped pool, but could not be sampled because of the riprap covering the bottom of the dredging area. Site water was collected from one sample location within the dredging area. A sufficient volume of sediment was collected for chemical analysis, physical analysis, standard and effluent elutriate generation, and TCLP preparation and testing. Upon completion of field activities, the sediment and water samples were submitted to TestAmerica for physical and chemical analysis. Sediment collection occurred on July 20 and 21, 2010, and site water collection occurred on July 21, 2010. Sediment samples were processed at EA's warehouse in Sparks, Maryland, on July 22, 2010.

The oyster survey sampling procedures are provided in Appendix A.

2.1 Sampling Locations

Sampling locations were chosen by EA in consultation with UniStar prior to the start of sampling. Coordinates for sediment samples (Maryland State Plane NAD83, feet) are provided in Table 1 and sampling locations are shown in Figure 2. Positioning was determined in the field using a differential global positioning system (DGPS).

2.2 In Situ Water Quality Measurements

Water quality measurements were recorded *in situ* at the surface, mid-depth, and bottom of the water column at each sampling location using a YSI water quality probe. The following parameters were measured (Table 2):

- Temperature
- pH
- Dissolved oxygen
- Conductivity
- Salinity
- Turbidity

Date and time, sampling location, and water depth were recorded at each sampling location.

2.3 Site Water and Elutriate Water

Approximately 24 gallons of site water and elutriate preparation water were collected from one location on July 21, 2010 (Figure 2). Water was collected from mid-depth of the water column using ISCO pumps with dedicated Tygon tubing. Site water for analytical testing was pumped directly into laboratory-prepared sample containers and shipped from the field on the day of collection via overnight delivery to TestAmerica in Pittsburgh, Pennsylvania. The elutriate preparation water was placed in 1-gallon certified cleaned, amber glass bottles, stored on ice in coolers after collection, and transported to EA's Sparks, Maryland, office where it was stored in

a walk-in cooler refrigerated at 4 degrees Celsius (°C). Elutriate preparation water was hand delivered with the sediment samples to TestAmerica. Holding times for water samples began when the water was collected and transferred into the appropriate sample containers. Sample containers, preservation techniques, and holding requirements for water samples for chemical analyses are provided in Table 3.

2.4 Sediment Sampling

Sediment samples were collected at five of the six target sediment sample locations in the proposed dredging area (Figure 2 and Table 1). These five locations represented two of the dredging areas: the barge unloading facility and the discharge pipe. Because of rip rap and rock on the sediment surface, samples could not be collected at the wedge shaped pool sample location. No sampling was conducted at the fish return dredging area because at the time sampling was conducted, material excavated from this area was expected to be returned to the area of excavation after installation of the fish return. This material will now be placed with the rest of the project dredged material. Sampling was conducted from Athena Technologies' 24-foot pontoon boat outfitted with a sampling platform and vibracoring system. A vibracorer outfitted with cellulose acetate butyrate (CAB) plastic liners (2.8-inch inner diameter) was used to collect the sediment samples. The stainless steel core barrel was fitted with a one-way valve at the top to retain sediment within the liner during retrieval. The core barrel was also fitted with a stainless steel nose cone to facilitate sediment penetration.

To obtain sediment cores, a dedicated, decontaminated core liner was loaded into the core barrel. The barrel was fitted with the nose cone and deployed through a moon pool in the center of the pontoon boat. The vibracore was lowered into the water until the nose cone was in contact with the sediment surface. The vibrating mechanism was started, causing the core barrel to shake, disturbing the sediment that is in contact with the sample barrel, reducing friction, and increasing the penetration of the core barrel into sediment.

After the desired sediment depth was reached or refusal occurred, the vibracorer was retrieved and brought aboard the pontoon boat. The core liner was removed from the steel barrel, and excess liner was cut to the sediment interface with a decontaminated hacksaw/blade. The core was capped at both ends, sealed, and labeled. Cores were kept on-board the boat on ice until the end of each work day and stored in a secured area at the landside staging area on ice until the sediment sampling was completed. At the completion of the sampling effort, the cores were transferred to a refrigeration unit cooled to 4°C at EA's office in Sparks, Maryland, and stored until processing. Core sample numbers, dates and times were recorded on a Chain-of-Custody (COC) form. The number of cores from each location, day and time collected, penetration depth, and recovery depth is reported in Table 1.

2.5 Sample Processing and Compositing

The sediment cores were processed in a designated area at EA's warehouse in Sparks, Maryland, on July 22, 2010. Prior to processing, cores were sorted and checked against the COC form. Multiple cores from each sampling location were composited and sub-sampled for physical and chemical analysis. Sediments were extracted from each core section using a stainless steel

extrusion rod and each section was homogenized in a stainless steel buckets or tubs until the sediment was thoroughly mixed and of uniform consistency. Sample processing equipment that came into direct contact with the sediment was decontaminated according to protocols specified in Section 2.6.

Composite samples were prepared for the samples collected in the vicinity of the barge unloading facility. The following compositing scheme was used:

Location	-	-	-	Sample ID
CCU3-Barge1	+	CCU3-Barge2	=	CCU3-BAR-1/2
CCU3-Barge3	+	CCU3-Barge4	=	CCU3-BAR-3/4

After the samples were homogenized and composited, samples of the sediment were removed for target analyses, placed into pre-cleaned glass jars using stainless steel spoons, and labeled. Sample containers, preservation techniques, and holding requirements for sediment and TCLP samples for chemical analyses are provided in Table 4. Holding times for the sediment samples began when the sediment was removed from the core liner, composited, homogenized, and placed in the appropriate sample containers.

2.6 Equipment Decontamination Procedures

Equipment that came into direct contact with sediment during sampling was decontaminated prior to deployment in the field to minimize cross-contamination. This included CAB core liners, core caps, stainless steel cutters, stainless steel catchers, and stainless steel processing equipment (spoons, knives, bowls, extruder, etc.). Any equipment that was reused in the field was decontaminated on-board the sampling boat between locations. While performing the decontamination procedure, phthalate-free nitrile gloves were used to prevent phthalate contamination of the sampling equipment or the samples.

The decontamination procedure is described below:

- Rinse equipment using clean tap or site water
- Rinse with 10 percent nitric acid (HNO₃)
- Rinse with distilled or de-ionized water
- Rinse with methanol followed by hexane
- Rinse with distilled or de-ionized water
- Air dry (in area not adjacent to the decontamination area)

Waste liquids produced during decontamination procedures were contained at the areas of decontamination. Decontamination waste liquid produced on-board the vessel was collected in 5-gallon buckets with lids and transferred to a 55-gallon secure drum at EA's warehouse in Sparks, Maryland, at the end of the project. Decontamination waste liquid generated at the sample processing area was contained directly in the secure drum at the warehouse. The liquid contained in the drums will be tested, characterized, and disposed of by a subcontractor.

2.7 Equipment Blank

Equipment blanks are collected to determine the extent of contamination, if any, from the sampling equipment used as part of the project. One equipment blank was collected on July 21, 2010. The equipment blank was collected by pouring de-ionized water through unused CAB liner from the supply that was used for sampling. The water was collected in appropriate containers and submitted for analysis.

The equipment blank was shipped via overnight delivery to TestAmerica on the day of collection. The sample containers, preservatives, and holding time requirements for the equipment blank are provided in Table 3. Holding times for the equipment blank began when the samples were collected and placed into the appropriate sample containers.

2.8 Documentation

Field notes were recorded in a permanently bound, dedicated field logbook. A log of coring activities, sampling locations, water depths, and core recoveries were recorded in the log. Coordinates and approximate water depth was recorded for each sampling location. Personnel names, local weather conditions, and other information that may impact the field sampling program were also recorded. Each page of the logbook was dated by the personnel entering the information. Copies of the logbooks were filed at EA's office in Sparks, Maryland, and are provided in Appendix B.

2.8.1 Numbering System

The sample numbering system was used to communicate between the field crew and the analytical laboratory, and indicates which samples were collected from each location.

Samples were labeled as follows:

Example:	CCU3-DIS	Sediment sample from one location
	CCU3-BAR-1/2	Sediment sample composite of multiple locations

The first two letters denoted the site designation (CCU3 = Calvert Cliffs), the next two digits denoted the project at the site (U3 = Unit 3), and the next three digits denoted the sampling area (DIS = discharge pipe area, BAR = barge dock area). For areas where multiple samples were collected from within the sampling area, one or more digits were added to identify the sampling location number. The sample identification was followed by one of the suffixes according to sample type:

- SED sediment sample to be submitted for chemical and physical analyses
- SW site water to be submitted for chemical analyses or elutriate preparation
- EET effluent elutriate sample
- SET standard elutriate sample

One sediment sample and one site water sample were designated for matrix spike (MS) / matrix spike duplicate (MSD) analysis by adding –MS or –MSD at the end of the sample name.

2.8.2 Sample Documentation

Both the individual sediment cores and the processed sediment samples were labeled. Sediment cores collected in the field were labeled with the sampling location number, core orientation (top and bottom), and date of collection. Sample containers for the processed sediment and water samples were labeled with the following information:

- Client name
- Project number
- Sample ID
- Sampling location
- Date and time of collection
- Sampler's initials
- Type of analyses required

2.8.3 Chain-of-Custody Records

Sediment, site water, and elutriate water samples were documented on a COC form. The COC form indicated the date and time of sample collection and was signed by the appropriate personnel. The COC form accompanied the samples to the analytical laboratory (Appendix C).

3. ANALYTICAL METHODS AND DATA EVALUATION

The majority of the analytical testing of site water, sediment, and elutriates for this characterization was conducted by TestAmerica – Pittsburgh, located in Pittsburgh, Pennsylvania. Additional services were provided by TestAmerica's laboratories in Burlington, Vermont (geotechnical parameters); North Canton, Ohio (total Kjeldahl nitrogen); and Knoxville, Tennessee (2,3,7,8-TCDD); and by a soil laboratory at Virginia Tech [calcium carbonate exchange (CCE), potential peroxide acidity (PPA), concentrated paste pH, and electrical conductivity.

3.1 Analytical Methods

All inorganic and organic compounds for these projects were determined using the methods listed in Table 5 as described in the laboratory's analytical SOPs. To meet program-specific regulatory requirements for chemicals of concern, all methods/SOPs were followed as stated with some specific requirements noted below:

Total Organic Carbon (TOC)

TOC in sediments was determined using the 1988 EPA Region II combustion oxidation procedure (the Lloyd Kahn procedure).

Polynuclear Aromatic Hydrocarbons – PAHs

To achieve the target detection limits (TDLs) referenced in QA/QC Guidance for Sampling and Analysis of Sediments, Water, and Tissues for Dredged Material Evaluations - Chemical Evaluations (EPA 823-B-95-001, April 1995), the PAHs were determined utilizing SW846 Method 8270C using Selective Ion Monitoring (SIM). For those samples where both semivolatiles by SW846 Method 8270C and PAHs by SW846 Method 8270C SIM were requested, both analyses were performed on the same extract. For those samples, the evaluation of method performance was based on the determined recoveries of surrogates and control analytes (in the LCS and MS/MSDs) from the semivolatiles by 8270C (full scan GC/MS) analyses because the spiked concentrations exceeded the calibration range for the PAH by GC/MS SIM analyses.

Standard Elutriate Test

The Standard Elutriate Test (SET) was used to predict the release of contaminants to the water column resulting from open water placement of dredged material. The SET was performed following the procedures in the *ITM* (USEPA/USACE 1998). For the SET, the laboratory creates the elutriate based on a sediment-to-water ratio of 1:4, on a volume basis. The sediment and site water volume requirements needed for the SET was dependent on the number and type of analytical tests to be performed on the elutriate.

Standard elutriates were prepared by using the site water collected onsite and sediment composites or individual sample locations (see Section 2.5). A sediment/water mixture was thoroughly mixed for 30 minutes. The mixture was then allowed to settle, and the supernatant was siphoned off, filtered to remove particulates, and then analyzed for the dissolved chemical constituents specified in Table 5. The reported results from the SET included a "dissolved"

value for each of the target parameters to be determined. Quantitation limits were the same as aqueous samples (Appendix D)

Effluent Elutriate Test

The Effluent Elutriate Test (EET) was used to predict the quality of effluent discharged from confined dredged material disposal area during dredging operations. The EET was performed following the procedures in the ITM, Appendix B, June 1998 (USEPA/USACE 1998). Effluent elutriates were prepared by using the site water collected onsite and sediment composites or individual sample locations (see Section 2.5).

The sediment and site water were mixed in a ratio equal to the average inflow concentration, and the mixture was manually mixed and aerated for one hour. That mixture was then allowed to settle for a time period equal to the anticipated field mean retention time, up to a period of 24 hours. The supernatant was then siphoned off and filtered to remove particulates, then analyzed for the dissolved chemical constituents specified in Table 5. For the EET, TestAmerica - Pittsburgh used the method default values of 120 g/L for the average field inflow concentration and 24 hours for the field mean retention time, respectively. The sediment and site water volume requirements needed for the EET was dependent on the number and type of analytical tests to be performed on the elutriate. Quantitation limits were the same as aqueous samples (Appendix D).

Toxicity Characteristic Leaching Procedure

TCLPs, which are routinely required for material placement at landfills and upland locations, are used to identify the potential for toxicity and to determine if the dredged material would be classified as a hazardous waste. TCLPs were prepared by using the site water from one location and sediment composites created from multiple locations (see Section 2.5). The sediment composites were extracted following the TCLP procedures specified in SW-846 Method 1311, and the resultant leachates were analyzed for the parameters specified in Table 5.

3.2 Detection Limits

Target detection limits (TDLs) for sediment, TCLP, and site water/elutriate samples are provided in Appendix D. The detection limit is a statistical concept that corresponds to the minimum concentration of an analyte above which the net analyte signal can be distinguished with a specified probability from the signal because of the noise inherent in the analytical system. The method detection limit concept (MDL) was developed by USEPA, and is defined as "the minimum concentration of a substance that can be measured and reported with 99 percent confidence that the analyte concentration is greater than zero" (40 CFR 136, Appendix B). Laboratory-specific target MDLs applicable to this project are provided in Appendix D. All analytical parameters, except wet chemistry parameters, butyltins, and 2,3,7,8-TCDD were quantitated to the MDL. All detected values quantified as greater than/equal to the MDL but less than the laboratory reporting limit (RL) were qualified as estimated. The laboratory RL is the lowest concentration at which an analyte can be detected in a sample and can be reported with a reasonable degree of accuracy and precision.

For sediment analyses, sample weight was adjusted for percent moisture (up to 50 percent moisture), prior to analysis, where appropriate to achieve the lowest possible reporting limits.

3.3 Laboratory Quality Control Samples

Internal laboratory quality assurance/quality control samples (including method blanks, laboratory control samples, surrogates, MS, and MSD) were analyzed.

- The **method (reagent) blank** is used to monitor laboratory contamination. The method blank is usually a sample of laboratory reagent water processed through the same analytical procedure as the sample (i.e., digested, extracted, distilled). One method blank was analyzed, at a frequency of one per every analytical preparation batch of 20 or fewer samples.
- The Laboratory Control Sample (LCS) is a fortified method blank consisting of reagent water or solid fortified with the analytes of interest for single-analyte methods or selected analytes for multi-analyte methods according to the appropriate analytical method. LCS's were prepared and analyzed with each analytical batch, and analyte recoveries are used to monitor analytical accuracy and precision.
- **Surrogates** are organic compounds that are similar to analytes of interest in chemical composition, extraction, and chromatography, but are not normally found in environmental samples. These compounds were spiked into all blanks, standards, samples, and spiked samples prior to analysis for organic parameters. Generally, surrogates are not used for inorganic analyses. Percent recoveries were calculated for each surrogate. Surrogates were spiked into samples according to the requirements of the reference analytical method. Surrogate spike recoveries were evaluated against the standard laboratory acceptance criteria limits, and were used to assess method performance and sample measurement bias. (If sample dilution caused the surrogate concentration to fall below the quantitation limit, surrogate recoveries were not calculated.)
- A **sample duplicate** is a second aliquot of a field sample that is analyzed to monitor analytical precision associated with that particular sample. Sample duplicates were performed for every batch of 20 or fewer samples.
- A MS is a field sample to which a known amount of analyte is added before sample preparation and analysis to evaluate the potential effects of matrix interference. Analyte concentrations in the spiked and unspiked sample were used to calculate percent recovery as a measure of matrix interference. A MSD is a duplicate of the MS sample.

3.4 Data Analysis

3.4.1 Calculations for Total PCBs and Total PAHs

For each sample, total PCB concentrations were determined by summing the concentrations of the 18 summation congeners [as specified in Table 9-3 of the Inland Testing Manual (ITM) (USEPA/USACE 1998)] and multiplying the total by a factor of two. Multiplying by a factor of

two estimated the total PCB concentration and accounted for additional congeners that were not tested as part of this program. These determinations were based upon testing of specific congeners recommended in the ITM and upon the National Oceanic and Atmospheric Administration (NOAA) (NOAA 1993) approach for total PCB determinations.

Total PAH concentrations were determined for each sample by summing the concentrations of the individual PAHs. To calculate the total PCB and total PAH concentrations, non-detects (ND) were assumed to be present at the MDL.

Substituting the MDL for non-detects (ND=MDL) provides a conservative estimate of the concentration. This method, however, tends to produce results that are biased high, especially in data sets where the majority of samples are non-detects. This overestimation is important to consider when comparing the calculated total values to criteria values.

3.4.2 Comparison to Sediment Quality Guidelines

Concentrations of detected analytes in sediment samples were compared to sediment quality guidelines (SQGs) for marine sediments, where available (Long et al. 1995). SQGs are tools which relate the concentrations of contaminants in sediment to predicted frequency or intensity of biological effects (Batley et al. 2005), and are intended to be either protective of biological resources or predictive of adverse effects to those resources, or both (Wenning and Ingersoll 2002). SQGs were developed as informal (non-regulatory) guidelines for use in interpreting chemical data from analyses of sediments. SQGs can be used to classify sediment samples with regard to their potential toxicity, to identify contaminants of concern, and to prioritize areas of concern based on the frequency and magnitude by which the values are exceeded (Long and MacDonald 1998).

Several biological-effects approaches have been used to assess marine/estuarine sediment quality relative to the potential for adverse effects on benthic organisms, including the Effects Range-Low (ERL) / Effects Range-Median (ERM) (Long et al. 1995) (Table 6). For dredged material evaluations, SQGs are used as a tool to assist with identification of constituents of potential concern (COPCs) and to provide additional weight of evidence in the evaluation (USACE–WES 1998).

Because they are based on environmental samples, ERLs and ERMs implicitly deal with contaminant mixtures, and the measured biological effects reflect the cumulative interactions of all chemicals in the mixture (Batley et al. 2005). The ERL and ERM values were derived by evaluating published measures of adverse biological effects, including field surveys of benthic and fish communities, Median Effective Concentrations (EC_{50}) and Lethal Concentration 50 (LC_{50}) values determined in laboratory bioassays, and toxicity predicted by equilibrium partitioning (Long et al. 1995). ERLs were established at the lower 10th percentile of the effects are unlikely (Long et al. 1995). ERMs were established at the lower 50th percentile of the effects data distribution, and represent concentrations above which adverse biological effects are probable (Long et al. 1995). Concentrations that are between the ERL and ERM represent the

concentrations at which adverse biological effects might occur. ERL and ERM benchmarks for marine/estuarine sediments are provided in Table 6.

3.4.3 Comparison to USEPA/State of Maryland Water Quality Criteria

Analytes detected in the site water and standard elutriate samples were compared to United States Environmental Protection Agency (USEPA) and State of Maryland saltwater acute and chronic water quality criteria for aquatic life (Table 7). Criteria were derived from USEPA's *National Recommended Water Quality Criteria (2010)* and the Code of Maryland Regulations (COMAR) (COMAR 26.08.02.03-2). The USEPA's acute criteria are based on 1-hour average exposure concentrations. The USEPA's chronic criteria are based on 4-day average exposure concentrations, with the exception of ammonia which is applied as a 30-day average exposure concentration.

The State of Maryland's saltwater acute and chronic water quality criteria for aquatic life are the same as the USEPA's. The USEPA and State of Maryland acute and chronic saltwater quality criteria for metals were developed for <u>dissolved</u> metal concentrations, but they are compared to total metals concentrations in this study as a conservative evaluation of the analytical results.

3.4.4 Calculation of Acute and Chronic Ammonia (NH₃-N) Criteria

The USEPA acute and chronic criteria for determining the toxicity of ammonia (NH₃-N) to aquatic life are variable, depending on temperature, pH, and salinity of the waterbody. The acute and chronic ammonia criteria for this analysis were based on the temperature, salinity, and pH measured at the site when the site water was collected (Table 2). The calculated acute ammonia criterion was 3.0 mg/L, and the calculated chronic ammonia criterion was 0.45 mg/L.

3.4.5 Evaluation of TCLP Data

To provide the information needed to determine if material could be placed in an upland beneficial use site or upland placement facility, concentrations of chemical constituents in the TCLP leachate were compared to maximum concentrations of contaminants for toxicity characteristics (Table 8). The toxicity characteristics are used to determine if a material should be classified as hazardous waste (40 CFR 261.24). The TCLP test is routinely required for dredged material placement at landfills and upland locations.

3.4.6 Comparison to Shirley Plantation Screening Criteria

Sediment results were also compared to values on Shirley Plantation's screening table (Table 9) to determine whether the sediment would be acceptable for placement at Shirley. This table is maintained by staff at Shirley Plantation in coordination with soil scientists at Virginia Tech.

4. ANALYTICAL RESULTS

The physical and chemical characteristics of sediment samples from the Calvert Cliffs dredging area were determined to assess the sediment quality of the material proposed for dredging. This chapter presents the results of the bulk sediment, site water, standard elutriate, and effluent elutriate chemical and TCLP analyses. The results are compared to the applicable guidelines or criteria.

Results of the sediment analyses were compared to sediment quality guidelines (ERLs/ERMs) and are shown in Tables 10 through 18. Results of the standard and effluent elutriate analyses and site water analyses are shown in Tables 19 through 26. Results of the TCLP analysis are presented in Table 39. Definitions of organic and inorganic data qualifiers are provided on the data results tables. Values for detected chemical constituents are shaded and bolded in the data tables. MDLs or RLs are presented for non-detected chemical constituents.

The results of the oyster survey are provided in Appendix A and summarized in Section 5.3.

4.1 Bulk Sediment Results

Sample weights were adjusted for percent moisture (up to 50 percent moisture) prior to analysis to achieve the lowest possible detection limits, and analytical results are reported on a dry weight basis. Copies of final raw data sheets (Form I's) and analytical narratives that include an evaluation of laboratory quality assurance/quality control results are available from EA upon request. The COC forms are provided in Appendix C.

4.1.1 Physical Analyses

Results of the grain size analyzes for sediment samples are shown in Table 16. The grain size analysis indicated that the Calvert Cliffs sediments are predominantly comprised of sand, ranging from 65.0 (CCU3-DIS-SED) to 82.1 percent (CCU3-BAR-3/4-SED) sand in the three sediment samples (Table 10).

4.1.2 General Chemistry Analyses

Results of the general chemistry analyses for sediment samples are shown in Table 11. Concentrations of total organic carbon (TOC) ranged from 0.17 percent to 1.51 percent.

Concentrations of ammonia as nitrogen ranged from 16.6 milligrams (mg)/kilogram (kg) to 137 mg/kg, nitrate-nitrite concentrations ranged from 0.61 to 5.5 mg/kg, and concentrations of total Kjeldahl nitrogen ranged from 319 to 749 mg/kg. Total phosphorus ranged from 215 to 841 mg/kg and total sulfide ranged from 39.4 to 354 mg/kg. Cyanide was not detected in the Calvert Cliffs sediment samples.

Two fractions of total petroleum hydrocarbons (TPH) were analyzed – diesel range organics (DRO) and gasoline range organics (GRO). TPH - DRO was detected in the two samples from the vicinity of the barge dock at concentrations of 40 and 370 μ g/kg (Table 11). TPH – GRO

was not detected in any of the sediment samples. There are no SQGs for TPH, but the Maryland Department of the Environment (MDE) has set generic cleanup standards for TPH for residential and non-residential soils. The residential cleanup standard (diesel+gasoline fractions) is 230,000 μ g/kg and the non-residential cleanup standard is 620,000 μ g/kg (MDE 2008). Concentrations of TPH-DRO detected in the Calvert Cliffs sediments were below the cleanup standard.

4.1.3 Inorganic Constituents

Of the 24 tested metals, 23 were detected in at least one of the sediment samples and the remaining metal (mercury) was detected in one of the three sediment samples (Table 12). No metals were detected in concentrations above the SQGs.

4.1.4 Organic Constituents

Results of the analysis of organic constituents in the Calvert Cliffs sediments are presented in Tables 13 to 18. Twelve PAHs, 16 PCB congeners, one PCB aroclor, five chlorinated pesticides, and two SVOCs were detected in the sediment samples. Only one constituent, total PCBs, exceed the SQGs. Total PCBs were detected at 1.25 times the ERL value, but at a concentration less than one fifth of the ERM value.

4.2 Site Water and Elutriate Results

Three effluent and three standard elutriates were prepared using the sediment samples. Chemical analyses for target analytes were conducted for each of the elutriate samples and the site water (Tables 19 through 26). Results of the analyses were compared to USEPA and State of Maryland acute and chronic saltwater criteria for the protection of aquatic life (Table 7). Values for detected constituents are shaded and bold in the data tables. Detection limits are presented for non-detected chemical constituents.

4.2.1 General Chemistry Analyses

Concentrations of general chemistry constituents in site water and elutriate samples are presented in Table 19. Chronic criteria exist for total sulfide and chronic and acute criteria ammonia. Acute and chronic ammonia criteria were calculated based on the temperature, pH, and the salinity in the water column at the time of site water collection. Ammonia concentrations exceeded the calculated chronic and acute criteria for the all the standard and effluent elutriate samples and the site water sample. Total sulfide was not detected in the effluent elutriate samples and dissolved sulfide was not detected in the standard elutriate samples.

TPH - DRO was detected in all of the site water, effluent elutriate, and standard elutriate samples. TPH - GRO was not detected in the site water, effluent elutriate, and standard elutriate samples. While there are no water quality standards for TPH, MDE has set a residential cleanup standard of 47 μ g/L for TPH (DRO + GRO fractions) in groundwater. All concentrations of TPH – DRO were above the residential cleanup standards for groundwater. The highest detected concentration of TPH – DRO was 230 μ g/L, which is 4.9 times the cleanup standard.

4.2.2 Inorganic Constituents

Concentrations of inorganic constituents tested in site water and elutriate samples are presented in Table 20. Eighteen of the 24 tested metals were detected in the site water sample. Concentrations of detected metals were low in site water, and did not exceed the acute or chronic water quality standards for aquatic life. Eighteen of the 24 tested metals were detected in the standard elutriate samples and 20 of the 24 tested metals were detected in the effluent elutriate samples. Detected concentrations of metals were generally low in all elutriate samples. Two of the constituents (copper, nickel) were detected at concentrations that exceeded water quality standards. Copper was detected in a concentration the two times the acute water quality criterion in one standard elutriate sample and 1.2 times the acute water quality standard in one effluent elutriate sample. Copper was detected in a concentration 1.2 times the chronic water quality criterion in one standard elutriate and at 1.5 and 1.03 times the chronic criterion in two effluent elutriate samples. Nickel was detected in a concentration 1.2 times the chronic criterion in one standard elutriate and at 1.5 and 1.03 times the chronic criterion in two effluent elutriate samples. Nickel was detected in a concentration 1.2 times the chronic criterion in one

4.2.3 Organic Constituents

Detected concentrations of organic constituents in site water and elutriate samples are presented in Tables 21 through 26. Six PAHs were detected in the site water sample, three PAHs were detected in the standard elutriates, and 15 PAHs were detected in the effluent elutriates. All detected concentrations in the site water and standard elutriates were below the reporting limit. Ten of the PAHs detected in the effluent elutriates were detected at concentrations below the reporting limit.

There were no PCB congeners or PCB aroclors detected in the site water sample. Four PCB congeners were detected in the standard elutriate samples and 1 PCB congener was detected in the effluent elutriate samples. Concentrations of detected PCBs were low in standard and effluent elutriate samples. PCB aroclors were not detected in the effluent and standard elutriate samples.

None of the tested chlorinated pesticides were detected in the site water sample. Only one of the tested chlorinated pesticides was detected in the standard elutriate samples. Five chlorinated pesticides were detected in the effluent elutriate samples. One chlorinated pesticide, heptachlor, was detected at concentrations up to 2.8 times the chronic criterion in the effluent elutriate samples.

Only one SVOC was detected in the site water sample and it was detected below the reporting limit. Three SVOCs were detected in the standard elutriate samples and one SVOC was detected in the effluent elutriate samples. There are no water quality criteria for the SVOCs detected in the standard and effluent elutriates.

Butyltins were only analyzed for the site water sample and were not detected. One dioxin congener (2,3,7,8-TCDD) was analyzed and was not detected in the site water or elutriate samples.

4.3 TCLP Results

The sediment composites were extracted following the TCLP procedures specified in SW-846 Method 1311. TCLPs were prepared using the sediment composites created from two sample locations within the dredging area. Results of the TCLP analysis are presented in Table 27. The samples were not ignitable and pH ranged from 7.9 to 8.8. Five of the metals (arsenic, barium, cadmium, mercury, and selenium) were detected at low concentrations that were below the TCLP screening value. None of the herbicides, pesticides, SVOCs, or volatile organic compounds (VOCs) were detected in the TCLP samples.

4.4 Shirley Plantation Screening

The Shirley Plantation Screening Table (Table 9) was populated with the results of the sediment sampling. The screening table instructions use the Proposed Virginia Clean Upland Fill Criteria and Exclusion Criteria to determine suitability for placement at the site. This table includes the results of four analyses used by staff at Shirley Plantation to make decisions related to the placement and management of the material onsite. These tests were to determine saturated paste pH, electrical conductivity, potential peroxide acidity, and calcium carbonate equivalency. Results are included in Table 28. None of the average concentrations of constituents exceed the Virginia Clean Upland Fill and Exclusion Criteria.

5. SUMMARY AND CONCLUSIONS

This study was conducted to characterize the material proposed for dredging offshore of CCNPP as part of the proposed Unit 3 project. This characterization includes determining whether or not the material would be suitable for placement at Shirley Plantation and whether or not a more detailed analysis of other placement options, such as beneficial use and innovative reuse, would be appropriate based on the quality of the material.

5.1 Physical Analyses

The results of the grain size analysis indicate that the material is predominantly sand, with concentrations of fine material (silts and clays) up to 26.7 percent. Grain size affects many beneficial use and innovative reuse projects. Examples of project types suitable for sandy material include: beach nourishment, island restoration, and wetland creation. Some options, such as oyster reef creation, are not viable for sandy material. The grain size of the material would not preclude confined disposal or landfill placement options.

5.2 Chemical Analyses

Only one constituent, total PCBs, was detected above the sediment quality guideline (ERL). The elutriate testing indicated that PCB congeners are tightly bound to the sediments and are not likely to be released into the water column during dredging or placement. The sediment data were integrated into the Shirley Plantation screening table and the mean concentrations of each constituent on the table was compared to the placement criteria. None of the results exceeded the Proposed Virginia Upland Clean Fill and Exclusion Criteria provided by staff at Shirley Plantation.

Three constituents (ammonia, copper, and nickel) exceed chronic water quality criteria in standard elutriate samples. Three constituents (ammonia, copper, and heptachlor) exceeded chronic water quality criteria in effluent elutriate samples. Copper concentrations exceed the acute water quality criteria in one standard elutriate sample and one effluent elutriate sample. Concentrations of nickel were above the chronic water quality criteria in one of the standard elutriate samples. Ammonia is a natural degradation product of organic matter in anoxic sediments which may be released to the water column during dredging and placement of material. However, ammonia concentrations would be expected to dissipate quickly in the water column to concentrations below the calculated criteria. Heptachlor only exceeded the chronic criterion in one of the effluent elutriate samples and was only slightly above the water quality criterion.

None of the TCLP analysis constituents exceeded maximum concentrations of contaminants for toxicity characteristics (40 CFR 261.24). Concentrations of detected constituents were well below the toxicity characteristic criteria, and the results also indicated that the materials were not corrosive or ignitable. The sediments within the dredging area would not be identified as hazardous waste per USEPA criteria.

Based on the results of the chemical analyses, there would be no anticipated restrictions on the use of the material for beneficial use, innovative reuse, confined disposal, upland placement, or placement in a landfill. Comparison to SQGs and water quality criteria indicated that the material would be suitable for use in restoration and habitat development activities, which tend to have the most stringent criteria for use. The material is suitable for placement at Shirley Plantation based on the screening criteria provided by staff at the site. The TCLP analysis indicated that the material is not hazardous and would be acceptable for landfill cover or upland placement.

5.3 Oyster Survey

The oyster survey results are presented in greater detail in Appendix A. The bottom type at the survey sites was either hard bottom (sandstone, rock or shell hash) or sand. Most of the bottom surveyed in the barge slip area could potentially serve as oyster habitat, but the discharge pipe area had a sand substrate, which is not considered oyster habitat. Shell coverage in the proposed dredging area was low, with only 21 percent of the sites sampled containing shell hash and at those sites, only a low to moderate amount of shell was found. Three live oysters and no dead oysters were collected during the survey and they were all found in a single grab in one of the samples collected furthest from shore.

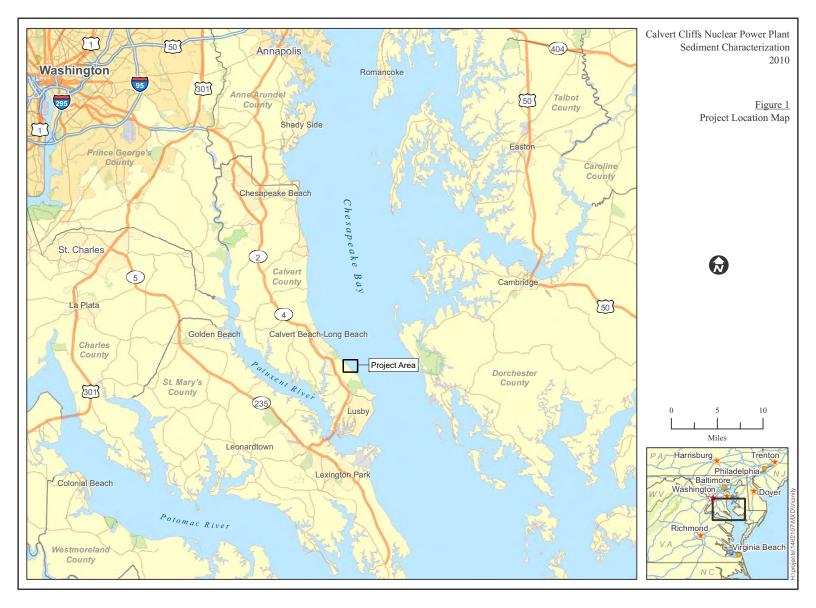
5.4 Conclusions

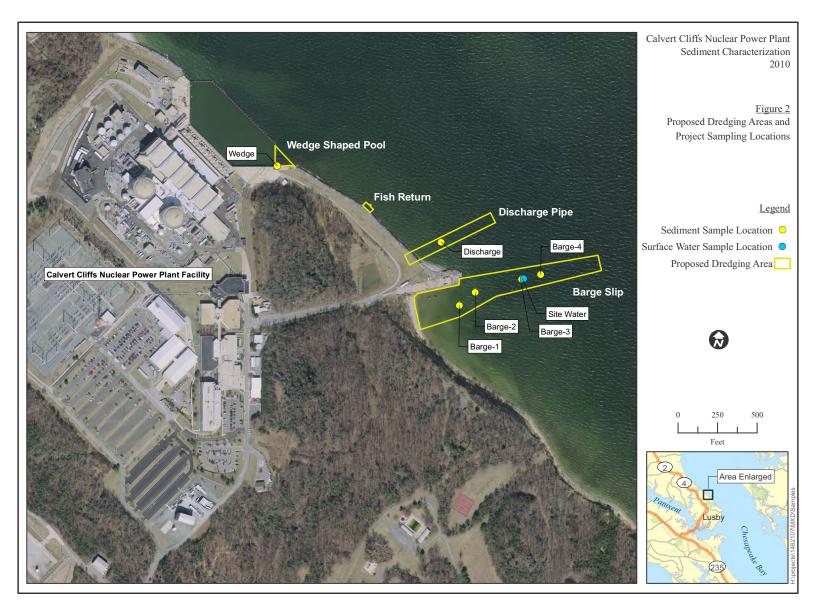
Based on the analyses conducted, the sediments proposed for dredging offshore of CCNPP in the Chesapeake Bay are composed predominantly of sand and would be viable, based on grain size, for use in a variety of beneficial use or innovative reuse projects. Although three constituents had concentrations that exceeded the sediment guidelines for one or more of the sediment samples, average concentrations of these constituents would be low and would likely be acceptable for any type of placement option. The grain size and quality of the CCNPP sediment would be acceptable for confined placement, beneficial use, innovative reuse, and landfill options. Sediment quality data were specifically compared to placement criteria for the Shirley Plantation placement site, which is an innovative reuse site where dredged material is being used for pit mine reclamation. CCNPP sediments would be acceptable for placement at Shirley Plantation. The sediment would be suitable for a range of beneficial use and innovative reuse options based on site specific placement criteria, available site capacity, and relative placement cost.

The results of the oyster survey indicate that the material dredged would is unlikely to have enough shell content to make recovery of oyster shell during dredging worthwhile. The amount of shell recovered would not be sufficient on its own for use for an oyster restoration project.

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Location	Northing (MD State Plane NAD 83, feet)	Easting (MD State Plane NAD 83, feet)	Date	Water Depth (feet)	Recovery (feet)	Sediment Volume of Recovery
Barge-1	279440.79	1474090.46	7/20/2010	5.33	7.5	2.4
Darge-1	279110.79	117 1090.10	//20/2010	5.55	8.33	2.7
					7.25	2.3
Barge-2	279525.34	1474191.54	7/20/2010	9.25	7.25	2.3
					6.08	1.9
					2.66	0.9
Barge-3	279610.35	1474482.89	7/20/2010	14.5	3.33	1.1
					3	1.0
	279639.62	1474605.49	7/20/2010		1.92	0.6
				15.83	1.75	0.6
Barge-4					1.42	0.5
Darge-4				15.65	1.17	0.4
					2.17	0.7
					2.5	0.8
Wedge	280347.59	1472953.99	7/20/2010	19.58		
weuge	280359.55	1472952.36	//20/2010	21.42		
	279842.88	1473973.72	7/20/2010	6.17	5.75	1.8
Disaharga	2/9042.00	14/39/3./2	7/20/2010	0.17	4.58	1.5
Discharge	279847.23	1473970.33	7/21/2010	7.42	4.08	1.3
	2/904/.23	14/39/0.33	7/21/2010	/.42	6.5	2.1
Surface Water Sample	279612.66	1474490.36	7/21/2010			

TABLE 1. SAMPLING LOCATIONS AND CORING SUMMARY
Calvert Cliffs Sediment Characterization (2010)

Location	Date & Time Sampled	Water Depth (ft)	Weather Conditions	Sample Depth	Water Temperature (°C)	pH	Dissolved Oxygen (mg/L)	Conductivity	Salinity (ppt)	Turbidity (NTU)					
			Clear, 75 to 95°C,	Surface	27.8	7.9	5.0	23.7	14.3	1.8					
Barge-1	7/20/10 0820	5.3	winds SW <5 knots	Middle	27.9	7.9	4.9	23.8	14.3	1.8					
			winds 5 w <5 kilots	Bottom	27.9	7.9	4.8	23.8	14.4	3.0					
			Class 75 to 05%	Surface	28.1	8.0	5.4	23.5	14.2	1.2					
Barge-2	7/20/10 0925	9.3	Clear, 75 to 95°C, winds SW <5 knots	Middle	28.1	8.0	5.3	23.6	14.2	1.1					
			winds 5 w <5 kilots	Bottom	28.1	7.9	5.1	23.7	14.3	1.6					
		14.5	GL 75 050G	Surface	28.7	8.1	6.4	23.6	14.2	1.7					
Barge-3	7/20/2010 1205		14.5	Clear, 75 to 95°C, winds SW <5 knots	Middle	28.6	8.0	5.9	23.8	14.3	1.4				
			winds SW <5 knots	Bottom	28.4	7.9	4.8	23.8	14.4	2.2					
		10 1310 15.8	/10 1310 15.8	/10 1310 15.8		G1 75 050G	Surface	28.9	8.1	6.9	23.3	14.0	2.2		
Barge-4	7/20/10 1310				.8 Clear, 75 to 95°C, winds SW <5 knots	Middle	28.6	8.0	5.8	23.6	14.2	1.5			
			winds S w <5 kilots	Bottom	28.3	8.0	5.3	23.6	14.2	1.7					
								G1 75 050G	Surface	28.4	7.9	5.4	23.8	14.3	1.6
Wedge	7/20/10 1114	19.6	Clear, 75 to 95°C, winds SW <5 knots	Middle	28.3	7.9	5.3	23.8	14.3	1.6					
			winds S w <5 kilots	Bottom	28.2	7.9	4.8	23.8	14.4	3.0					
			CI 75 (050C	Surface	29.5	8.4	8.0	23.4	14.1	3.6					
Discharge	7/20/10 1535	6.2	Clear, 75 to 95°C, winds SW <5 knots	Middle	29.2	8.4	8.3	22.9	13.8	2.5					
			winds 5 w <5 kilots	Bottom	29.2	8.4	8.4	22.9	13.7	1.9					
Site Water			Overcast, 70 to	Surface	28.1	8.1	6.4	22.8	13.7	1.4					
	7/21/10 1100	15.0	95°C, winds S <5	Middle	27.7	8.0	5.6	22.9	13.8	1.1					
(Barge 3)			knots	Bottom	28.3	7.9	4.5	23.5	14.1	1.0					

TABLE 2. IN SITU WATER QUALITY DATACalvert Cliffs Sediment Characterization (2010)

TABLE 3. REQUIRED CONTAINERS, PRESERVATION TECHNIQUES, AND HOLDING
TIMES FOR AQUEOUS SAMPLES ^(a)Calvert Cliffs Sediment Characterization (2010)

Parameter	Volume Required ^(b)	Container ^(c)	Preservative	Holding Time
Inorganics				
Metals (including Mercury)	1 Liter	Р	pH <2 with HNO ₃ Cool, 4°C	6 months (28 days for Hg)
Sulfide	500 mLs	P,G	NaOH to pH >9 Zinc Acetate Cool, 4°C	7 days
Nitrogen (Ammonia, Nitrate + Nitrite)	250 mLs	P,G	H ₂ SO ₄ to pH <2 Cool, 4°C	28 days
Nitrogen (Total Kjeldahl), Total Phosphorus	500 mLs	P,G	H ₂ SO ₄ to pH <2 Cool, 4°C	28 days
Physical Parameters				
Site Water for Standard Elutriate Test	5 gallons	G	Cool, 4°C	None specified
Site Water for Effluent Elutriate Test	5 gallons	G	Cool, 4°C	None specified
Organics				
Total Petroleum Hydrocarbons (TPH) Gas Range Organics (GRO)	2-40 mL	G, teflon-lined, septa cap	HCl, Cool, 4°C	14 days
Total Petroleum Hydrocarbons (TPH) Diesel Range Organics (DRO)	2 liters	G, teflon-lined cap	Cool, 4°C	7 days until extraction, 40 days after extraction
Total Organic Carbon	3-40 mLs	G, teflon-lined, septa cap	H ₂ SO ₄ or HCl to pH <2; Cool, 4°C	28 days
Chlorinated Pesticides, Polynuclear Aromatic Hydrocarbons, PCB Congeners, PCB Aroclors, Semivolatile Organic Compounds	6 Liters	G, teflon-lined cap	Cool, 4°C	7 days until extraction, 40 days after extraction
Dioxins (2, 3, 7, 8 TCDD only) / Furans	2-1Liters	G	4°C	7 days until extraction, 40 days after extraction

(a) From time of sample collection.

(b) Additional volume will need to be provided for samples designated as MS/MSDs

(c) P = plastic; G = glass.

TABLE 4. REQUIRED CONTAINERS, PRESERVATION TECHNIQUES, AND HOLDING TIMES
FOR SEDIMENT SAMPLES ^(a)
Calvert Cliffs Sediment Characterization (2010)

Parameter	Volume Required ^(b)	Container ^(c)	Preservative	Holding Time	
Inorganics					
Metals (including Mercury)	4 oz.	P,G	4°C	6 months (28 days for Hg)	
Nitrogen (Ammonia, Nitrate + Nitrite)	4 oz	P,G	4°C	28 days	
Nitrogen (Total Kjeldahl), Total Phosphorus	4 oz	P,G	4°C	28 days	
Sulfide	4 oz	P,G	4°C	7 days	
Physical Parameters					
Grain Size and Percent Solids	32 oz	P,G	4°C	6 months	
Sediment for Standard Elutriate Test	1-2 gallons	G	4°C	14 days	
Sediment for Effluent Elutriate Test	3-4 gallons	G	4°C	14 days	
Sediment for TCLP	4-4 oz 1-32oz	G	4°C	14 days to TCLP extraction, 7 days after extraction	
Sediment for Virginia Tech Analyses	1-8 oz	G	4°C		
Organics					
Total Organic Carbon	4 oz	G	4°C	14 days	
PAHs, PCB Congeners, Chlorinated Pesticides, Semivolatile Organic Compunds (SVOC)	32 oz	G	4°C	14 days until extraction, 40 days from extraction to analysis	
Butyltins	8 oz	G	4°C	14 days until extraction, 40 days from extraction to analysis	
Dioxins (2, 3, 7, 8 TCDD only)	4 oz	G	4°C	1 year until extraction, 50 days after extraction	
Total Petroleum Hydrocarbons (TPH) Diesel Range Organics (DRO)	4 oz.	G	4°C	14 days until extraction, 40 days after extraction	
Total Petroleum Hydrocarbons (TPH) Gas Range Organics (GRO)	4 oz.	G	4°C	14 days until analysis	

(a) From time of sample collection.

(b) Additional volume will need to be provided for samples designated as MS/MSD/MDs.

(c) P = plastic; G = glass.

Analyte	Analytical Method
Sediment and Elutriates	•
Metals (ITM list)	SW846 6020
Mercury	SW846 7471A
Polynuclear Aromatic Hydrocarbons (PAHs)	SW846 8270C SIM
Polychlorinated Biphenyl (PCB Congeners)	SW846 8082
Polychlorinated Biphenyl (PCB Aroclors)	SW846 8082
Chlorinated Pesticides	SW846 8081A
2,3,7,8-TCDD	EPA-5 1613B
Ammonia	EPA 350.1
Nitrate+Nitrite	EPA 353.2
Total Kjeldahl Nitrogen	EPA 351.2
Total Phosphorus	EPA 365.2
Total Sulfides	SW846 9034
Total Organic Carbon	Lloyd Khan / 415.1
Grain Size (sediment only)	ASTM D422
Percent Solids (sediment only)	SM 2540B
Total Petroleum Hydrocarbons (THP) Gas Range Organics (GRO)	SW846 8015
Total Petroleum Hydrocarbons (THP) Diesel Range Organics (DRO)	SW846 8015
Semivolatile Organic Compounds (SVOCs)	SW846 8270C LL
Calcium Carbonate Equivalence (CCE) (sediment only)	Virginia Tech Soil Lab – Sediment Only
Potential Peroxide Acidity (PPA) (sediment only)	Virginia Tech Soil Lab – Sediment Only
Saturated paste pH	Virginia Tech Soil Lab –
(sediment only) Electrical Conductivity (EC)	Sediment Only Virginia Tech Soil Lab –
(sediment only)	Sediment Only
Toxicity Characteristic Leaching Procedure (7	ГСLР) (sediment only)
TCLP Volatiles	SW846 8260B
TCLP Pesticides	SW846 8081A
TCLP Semivolatiles	SW846 8270C
TCLP Herbicides	SW846 8151A
TCLP Metals	SW846 6010B
TCLP Mercury	SW846 7471A
Ignitability	SW846 1010
Leaching Procedure	EPA 1311

TABLE 5. ANALYTICAL METHODSCalvert Cliffs Sediment Characterization (2010)

Constituent	Units	Effects Range Low (ERL)*	Effects Range Median (ERM)*
METALS			
ARSENIC	MG/KG	8.2	70
CADMIUM	MG/KG	1.2	9.6
CHROMIUM	MG/KG	81	370
COPPER	MG/KG	34	270
LEAD	MG/KG	46.7	218
MERCURY	MG/KG	0.15	0.71
NICKEL	MG/KG	20.9	51.6
SILVER	MG/KG	1	3.7
ZINC	MG/KG	150	410
CHLORINATED PESTICIDES	1	1	Г
CHLORDANE	UG/KG	0.5	6
4,4-DDD	UG/KG	2	20
4,4-DDE	UG/KG	2.2	27
4,4-DDT	UG/KG	1	7
DIELDRIN	UG/KG	0.02	8
GAMMA-BHC (LINDANE)	UG/KG		
PAHs	-	1	
2-METHYLNAPHTHALENE	UG/KG	70	670
ACENAPHTHENE	UG/KG	16	500
ACENAPHTHYLENE	UG/KG	44	640
ANTHRACENE	UG/KG	85.3	1,100
BENZO[A]ANTHRACENE	UG/KG	261	1,600
BENZO(A)PYRENE	UG/KG	430	1,600
CHRYSENE	UG/KG	384	2,800
DIBENZ(A,H)ANTHRACENE	UG/KG	63.4	260
FLUORANTHENE	UG/KG	600	5,100
FLUORENE	UG/KG	19	540
NAPHTHALENE	UG/KG	160	2,100
PHENANTHRENE	UG/KG	240	1,500
PYRENE	UG/KG	665	2,600
PAHs, TOTAL	UG/KG	4,022	44,792
PCBs			
PCBs, TOTAL	UG/KG	22.7	180
SEMIVOLATILE ORGANIC COMPO			
BIS(2-ETHYLHEXYL) PHTHLATE	UG/KG		

TABLE 6. MARINE SEDIMENT BENCHMARKSCalvert Cliffs Sediment Characterization (2010)

*Sources: Long et al. 1995 and MacDonald et al. 1996

		SALTWATER CRITERIA ⁽¹⁾				
		USEPA/MARYLAND	USEPA/MARYLAND			
ANALYTE	UNITS	ACUTE ^a	CHRONIC^b			
NUTRIENTS						
AMMONIA	MG/L	0.37 °	0.06 ^c			
DISSOLVED SULFIDE	MG/L		0.002			
TOTAL SULFIDE	MG/L		0.002			
METALS**						
ARSENIC	UG/L	69 ^e	36 ^e			
CADMIUM	UG/L	40 ^e	8.8 ^e			
CHROMIUM	UG/L	1,100 ^e	50 ^e			
COPPER ⁽²⁾	UG/L	4.8 ^e / 6.1	3.1 ^e			
LEAD	UG/L	210 ^e	8.1 ^e			
MERCURY	UG/L	1.8 ^e	0.94 ^e			
NICKEL	UG/L	74 ^e	8.2 ^e			
SELENIUM	UG/L	290 ^e	71 ^e			
SILVER	UG/L	1.9 ^{e,f}				
ZINC	UG/L	90 ^e	81 ^e			
CHLORINATED PESTICIDES						
4,4'-DDT	UG/L	0.13 ^f	0.001 ^f			
ALDRIN	UG/L	1.3 ^f				
CHLORDANE (TECHNICAL)	UG/L	0.09 ^f	0.004 ^f			
DIELDRIN	UG/L	0.71 ^f	0.0019 ^f			
ENDOSULFAN I	UG/L	0.034 ^{f,g}	$0.0087 \ {}^{\mathrm{f,g}}$			
ENDOSULFAN II	UG/L	0.034 ^{f,g}	$0.0087 {}^{ m f,g}$			
ENDRIN	UG/L	0.037 ^f	0.0023 ^f			
GAMMA-BHC (LINDANE)	UG/L	0.16 ^f				
HEPTACHLOR	UG/L	0.053 ^f	0.0036 ^f			
HEPTACHLOR EPOXIDE	UG/L	0.053 ^f	0.0036 ^f			
METHOXYCHLOR	UG/L		0.03			
MIREX	UG/L		0.001			
TOXAPHENE	UG/L	0.21	0.002			

TABLE 7. USEPA AND STATE OF MARYLAND ACUTE AND CHRONIC SALTWATER QUALITY CRITERIA FOR AQUATIC LIFE* Calvert Cliffs Sediment Characterization (2010)

*Source : USEPA 2008. National Recommended Water Quality Criteria; Code of Maryland Regulations (COMAR 26.08.02.03-2)

**Water quality criteria for the metals are based on dissolved concentrations.

(1) The State of Maryland's saltwater acute and chronic water quality criteria for the protection of aquatic life are equivalent to the USEPA criteria for each tested analyte

(2) The State of Maryland has an estuarine acute copper criterion of 6.1 μ g/L that applies to waters in Baltimore Harbor and the Upper Chesapeake Bay

Superscripts:

a = acute aquatic life criteria based on 1-hour average exposure concentrations

b = chronic aquatic life criterion based on 4-day average exposure concentrations

c = total ammonia as nitrogen, calculated based on site specific conditions

d = free cyanide as mg CN/L

e = saltwater criteria for metals are expressed in terms of the dissolved metal in the water column

f = instantaneous maximum

g = value for endosulfan I + endosulfan II

TABLE 8. TCLP REGULATORY GUIDELINESCalvert Cliffs Sediment Characterization (2010)

(Maximum Concentration of Contaminants for Toxicity Characteristics)

CHEMICAL NAME	REGULATORY LEVEL (MG/L)
METALS	
ARSENIC	5
BARIUM	100
CADMIUM	1
CHROMIUM	5
LEAD	5
MERCURY	0.2
SELENIUM	1
SILVER	5
PESTICIDES AND HERBICIDES	
2, 4, 5-TP (SILVEX)	1
2, 4-D	10
CHLORDANE	0.03
ENDRIN	0.02
HEPTACHLOR (AND ITS EPOXIDE)	0.008
GAMMA-BHC (LINDANE)	0.4
METHOXYCHLOR	10
TOXAPHENE	0.5
SEMIVOLATILE ORGANIC COMPOU	
o-CRESOL*	200
m-CRESOL*	200
p-CRESOL*	200
CRESOL	200
1, 4 DICHLOROBENZENE	7.5
2,4 DINITROTOLUENE	0.13
HEXACHLOROBENZENE	0.13
HEXACHLOROBUTADIENE	0.5
HEXACHLOROETHANE	3
NITROBENZENE	2
PENTACHLOROPHENOL	100
2, 4, 5-TRICHOROPHENOL	400
2, 4, 6-TRICHOROPHENOL	2
PYRIDINE	5
VOLATILE ORGANIC COMPOUNDS (VOCs)
BENZENE	0.5
CARBON TETRACHLORIDE	0.5
CHLOROBENZENE	100
CHLOROFORM	6
1, 2 DICHLOROETHANE	0.5
1, 1 DICHLOROETHYLENE	0.7
2-BUTANONE	200
TETRACHLOROETHYLENE	0.7
TRICHLOROETHYLENE	0.5
VINYL CHLORIDE	0.2
*If a m n Cresol concentration cannot be	

*If o-, m-, p-Cresol concentration cannot be differentiated, the total cresol concentration is used. The regulatory level of total cresol is 200 mg/L.

Source: 40 CFR 261.24 (1993)

TABLE 9. SHIRLEY PLANTATION SCREENING TABLE Calvert Cliffs Sediment Characterization (2010)

	rom yo yses g	our sample		Criteria						
	yses g	o nere								
	es 1 an direc	d 2 below tions		NJDEP (1997) Residential Soil Cleanup Criteria ³		Screening Levels , 2008) ⁴	EPA Part 503 Biosolids	USGS soil background metals ⁵	Proposed VA Exclusion Criteria ⁶	Proposed VA Clean Upland Fill Criteria
Sample ID	Date	Average Value ^{1,2}	PARAMETER		Industrial Soil	Residential Soil	Exceptional Quality	VA background metal levels		
CCU3	Jul-10	2,023	Metals (mg kg ⁻¹) Aluminum	NA	990,000	77,000			NA	NA
CCU3	Jul-10	0.07	Antimony	14	410	31			410	14
CCU3 CCU3	Jul-10 Jul-10	2.3	Arsenic Barium	20 700	1.6 19,000	0.39 15,000	41	5 244	41 19,000	20 700
CCU3	Jul-10	0.15	Beryllium	1	2,000	160		<1	2,000	160
CCU3 CCU3	Jul-10	0.35	Cadmium	39	810	70	39	<0.1	810	39
CCU3	Jul-10 Jul-10	10.6	Calcium Chromium	NA NA	NA 200	NA 39		23	NA 1,200	NA 200
CCU3	Jul-10	1.17	Cobalt	NA	300	23			300	NA
CCU3 CCU3	Jul-10 Jul-10	2.8	Copper Iron	600 NA	41,000 720,000	3,100 55,000	1,500	9	4,300 150,000	1,500 150,000
CCU3	Jul-10 Jul-10	3.7	Lead	400	800	400	300	26	800	300
CCU3	Jul-10	1,787	Magnesium	NA	NA	NA		265	NA	NA
CCU3 CCU3	Jul-10 Jul-10	67.0 0.02	Manganese Mercury	NA 14	NA 100	NA 7.8	17	295 0.06	NA 100	NA 14
CCU3	Jul-10	5.6	Nickel	250	69,000	14,000	420	9	1,000	250
CCU3 CCU3	Jul-10 Jul-10	597 0.36	Potassium Selenium	NA 63	NA 5,100	NA 390	100		NA 5 100	NA 63
CCU3 CCU3	Jul-10 Jul-10	0.36	Silver	63	5,100	390	100		5,100 5,100	63 110
CCU3	Jul-10	2,003	Sodium	NA	NA	NA			NA	NA
CCU3 CCU3	Jul-10 Jul-10	0.16 7.4	Thallium Vanadium	2 370	NA 5,200	NA 390			10 5,200	2 370
CCU3	Jul-10	18	Zinc	1,500	310,000	23,000	2,800	41	7,500	1,500
CCU3	Jul-10	0.34	Cyanide, Total	1,100	20,000	1,600			20,000	1,100
CCU3	Jul-10	0.00083	PCBS (mg kg ⁻¹) Aroclor 1016	NA	21	3.9			21	NA
CCU3	Jul-10	0.00108	Aroclor 1221	NA	0.62	0.17			0.62	NA
CCU3	Jul-10	0.00097	Aroclor 1232	NA	0.62	0.17			0.62	NA
CCU3 CCU3	Jul-10 Jul-10	0.00092 0.00053	Aroclor 1242 Aroclor 1248	NA NA	0.74 0.74	0.22 0.22			0.74 0.74	NA NA
CCU3	Jul-10	0.00783	Aroclor 1254	NA	0.74	0.22			0.74	NA
CCU3	Jul-10	0.00082	Aroclor 1260	NA	0.74	0.22			0.74	NA
CCU3	Jul-10	0.01298	Total Aroclor ⁸ Pesticides (mg kg ⁻¹)	0.49	25.2	5.1			25.2	0.49
CCU3	Jul-10	0.00008	4,4'-DDD	3	7.2	2			7.2	3
CCU3 CCU3	Jul-10 Jul-10	0.00009	4,4'-DDE 4,4'-DDT	2 2	5.1 7	1.4 1.7			5.1 7	2 2
CCU3	Jul-10 Jul-10	0.00010	Aldrin	0.04	0.11	0.029			0.11	0.04
CCU3	Jul-10	0.00015	alpha-BHC	NA	NA	NA			NA	NA
CCU3 CCU3	Jul-10 Jul-10	0.00015 0.00011	beta-BHC alpha-Chlordane	NA NA	NA NA	NA NA			NA NA	NA NA
CCU3	Jul-10	0.00011	gamma-Chlordane	NA	NA	NA			NA	NA
CCU3	Jul-10	0.00009	delta-BHC	NA 0.042	NA 0.11	NA 0.03			NA	NA
CCU3 CCU3	Jul-10 Jul-10	0.00010 0.00011	Dieldrin Endosulfan I	0.042 NA	3,700	370			0.11 3,700	0.042 NA
CCU3	Jul-10	0.00036	Endosulfan II	NA	3,700	370			3,700	NA
CCU3 CCU3	Jul-10 Jul-10	0.00006	Endosulfan Sulfate Endrin	NA 17	3,700 180	370 18			3,700 180	NA 17
CCU3	Jul-10	0.00011	Endrin aldehyde	NA	NA	NA			NA	NA
CCU3	Jul-10	0.00009	Endrin ketone	NA 0.52	NA	NA			NA 0.52	NA 0.52
CCU3 CCU3	Jul-10 Jul-10	0.00041 0.00034	gamma-BHC (Lindane) Heptachlor	0.52 0.15	NA 0.38	NA 0.11			0.52 0.38	0.52 0.15
CCU3	Jul-10	0.00011	Heptachlor epoxide	NA	0.19	0.053			0.19	NA
CCU3 CCU3	Jul-10 Jul-10	0.00012 0.00377	Methoxychlor Toxaphene	280 0.1	3,100 1.6	310 0.44			3,100 1.6	280 0.1
	5ai-10	0.000//	Semivolatiles (mg kg ⁻¹)	0.1	1.0	0.77			1.0	
CCU3	Jul-10	0.0053	Acenaphthene	3,400	33,000	3,400			33,000	3,400
CCU3 CCU3	Jul-10 Jul-10	0.0020 0.0037	Acenaphthylene Anthracene	NA 10,000	NA 170,000	NA 17,000			NA 170,000	NA 10,000
CCU3	Jul-10	0.0055	Benzo(a)anthracene	0.9	2.1	0.15			2.1	0.9
CCU3 CCU3	Jul-10 Jul-10	0.0064	Benzo(b)fluoranthene	0.9 0.9	2.1 21	0.15 1.5			2.1 21	0.9
CCU3 CCU3	Jul-10 Jul-10	0.0047	Benzo(k)fluoranthene Benzo(ghi)perylene	0.9 NA	21 NA	1.5 NA			21 NA	0.9 NA
CCU3	Jul-10	0.0011	Benzo(a)pyrene	0.66	0.21	0.015			0.66	0.21
CCU3 CCU3	Jul-10 Jul-10	0.0037 0.0015	bis(2-Chloroethoxy)methane bis(2-Chloroethyl) ether	NA 0.66	1800 0.9	180 0.19			1,800 0.9	NA 0.66
CCU3	Jul-10 Jul-10	0.028	bis(2-Ethylhexyl) phthalate	49	120	35			120	49
CCU3	Jul-10	0.0049	4-Bromophenyl phenyl ether	NA	NA	NA			NA	NA
CCU3	Jul-10 Jul-10	0.0077	Butyl benzyl phthalate Carbazole	1,100 NA	910 NA	260 NA			1,100 NA	910 NA
						NA			230	230
CCU3 CCU3	Jul-10	0.0045	4-Chloroaniline	230	NA					
	Jul-10 Jul-10 Jul-10	0.0045 0.0052 0.0012	4-Chloroaniline 4-Chloro-3-methylphenol 2-Chloronaphthalene	230 10,000 NA	NA NA NA	NA NA			10,000 NA	10,000 NA

Valere f					itts Sediment		()			
		our sample					Criteria			
ana	lyses g	o here								
6		101.1		NJDEP (1997)						
		d 2 below		Residential Soil		Screening Levels	EPA Part 503	USGS soil	Proposed VA	Proposed VA Clean
for	direc	tions		Cleanup Criteria ³	(EPA	, 2008) ⁴	Biosolids	background metals ⁵	Exclusion Criteria ⁶	Upland Fill Criteria ⁷
		Average			Industrial	Residential		VA background		
Sample ID	Date	Value ^{1,2}	PARAMETER		Soil	Soil	Exceptional Quality	metal levels		
CCU3	Jul-10	0.0047	2-Chlorophenol	280	5,100	390	· · · ·		5,100	280
CCU3	Jul-10	0.0065	4-Chlorophenyl phenyl ether	NA	NA	NA			NA	NA
CCU3 CCU3	Jul-10 Jul-10	0.0058 0.0013	Chrysene Dibong(a b)onthrocomo	9 0.66	210 0.21	15 0.015			210 0.66	9 0.21
CCU3	Jul-10 Jul-10	0.0015	Dibenz(a,h)anthracene Dibenzofuran	NA	NA	NA			NA	NA
CCU3	Jul-10	0.0070	Di-n-butyl phthalate	5,700	NA	NA			5,700	5,700
CCU3	Jul-10	0.0060	1,2-Dichlorobenzene	5,100	10,000	2,000			10,000	5,100
CCU3	Jul-10	0.0044	1,3-Dichlorobenzene	5,100	NA	NA			5,100	5,100
CCU3 CCU3	Jul-10 Jul-10	0.0041 0.0060	1,4-Dichlorobenzene 3,3'-Dichlorobenzidine	570 2	13 3.8	2.6 1.1			570 3.8	13 2
CCU3	Jul-10 Jul-10	0.0000	2,4-Dichlorophenol	170	1,800	1.1			5.8 1,800	170
CCU3	Jul-10	0.0060	Diethyl phthalate	10,000	490,000	49,000			490,000	10,000
CCU3	Jul-10	0.0087	2,4-Dimethylphenol	1,100	12,000	1,200			12,000	1,100
CCU3	Jul-10	0.0060	Dimethyl phthalate	10,000	NA	NA			10,000	10,000
CCU3 CCU3	Jul-10 Jul-10	0.0060 0.023	Di-n-octyl phthalate 4,6-Dinitro-2-methylphenol	1,100 NA	NA NA	NA NA			1,100 NA	1,100 NA
CCU3	Jul-10	0.068	2,4-Dinitrophenol	110	1,200	120			1,200	110
CCU3	Jul-10	0.0047	2,4-Dinitrotoluene	NA	1,200	120			1,200	NA
CCU3	Jul-10	0.0058	2,6-Dinitrotoluene	1	620	61			620	61
CCU3 CCU3	Jul-10 Jul-10	0.015 0.0063	Fluoranthene	2,300 2,300	22,000 22,000	2,300 2,300			22,000	2,300
CCU3	Jul-10 Jul-10	0.0063	Fluorene Hexachlorobenzene	2,500	1.1	0.3			22,000 1.1	2,300 0.66
CCU3	Jul-10	0.0013	Hexachlorobutadiene	1	22	6.2			22	1
CCU3	Jul-10	0.0060	Hexachlorocyclopentadiene	400	3,700	370			3,700	400
CCU3	Jul-10	0.0041	Hexachloroethane	6	120	35			120	6
CCU3 CCU3	Jul-10 Jul-10	0.0012 0.0043	Indeno(1,2,3-cd)pyrene Isophorone	0.9 1,100	2.1 1,800	0.15 510			2.1 1,800	0.9 1,100
CCU3	Jul-10 Jul-10	0.0043	2-Methylnaphthalene	NA	4,100	310			4,100	NA
CCU3	Jul-10	0.0040	2-Methylphenol	2,800	NA	NA			2,800	2,800
CCU3	Jul-10	0.0055	4-Methylphenol	2,800	NA	NA			2,800	2,800
CCU3	Jul-10	0.0010	Naphthalene	230	20	3.9			230	20
CCU3 CCU3	Jul-10 Jul-10	0.026 0.023	2-Nitroaniline 3-Nitroaniline	NA NA	NA 82	NA 18			NA 82	NA NA
CCU3	Jul-10 Jul-10	0.023	4-Nitroaniline	NA	82	23			82	NA
CCU3	Jul-10	0.0048	Nitrobenzene	28	280	31			280	28
CCU3	Jul-10	0.006	2-Nitrophenol	NA	NA	NA			NA	NA
CCU3	Jul-10	0.020	4-Nitrophenol	NA 0.66	NA 0.25	NA 0.060			NA	NA 0.25
CCU3 CCU3	Jul-10 Jul-10	0.0014 0.0052	N-Nitroso-di-N-propylamine N-Nitrosodiphenylamine	0.66 140	0.25 350	0.069 99			0.66 350	0.25 140
CCU3	Jul-10	0.0052	Pentachlorophenol	6	9	3			9	6
CCU3	Jul-10	0.016	Phenanthrene	NA	NA	NA			NA	NA
CCU3	Jul-10	0.029	Phenol	10,000	180,000	18,000			180,000	10,000
CCU3 CCU3	Jul-10 Jul-10	0.013 0.0031	Pyrene 1,2,4-Trichlorobenzene	1,700 68	17,000 400	1,700 87			17,000 400	1,700 68
CCU3	Jul-10 Jul-10	0.0031	2,4,5-Trichlorophenol	5,600	400 62,000	6,100			400 62,000	5,600
CCU3	Jul-10	0.0087	2,4,6-Trichlorophenol	62	160	44			160	62
			Dioxin and Furans (ng kg ⁻¹)							
CCU3	Jul-10	0.13	2,3,7,8-TCDD	NA	18	4.3			18	4.3
			Tributyltin (mg kg ⁻¹)					r	r	1
CCU3	Jul-10	0.0063	Tributyltin Compounds	L	180	18		L		L
CCU3	40381	0.16	Petroleum (mg kg ⁻¹) TPH-DRO	r				r		1
1005	40301	0.10	1111-0100							1

TABLE 9. SHIRLEY PLANTATION SCREENING TABLE Calvert Cliffs Sediment Characterization (2010)

Values from your sample analyses go here			Additional Analyses ⁹	Units and Reporting convention	Method	Proposed VA Exclusion Criteria ⁶	Proposed VA Clean Fill Criteria ⁷
Sample ID	Date	Average Value ^{1,2}					
CCU3	Jul-10	None Detected	Acid-Base Accounting (all samples > 0.25% total S) or H2O2 Potential Acidity	Tons CCE acid demand per 1000 Tons Material	EPA 600-2-78-054	-15 unless under water table	-5
			Pyritic S	% or g kg ⁻¹		2.00	0.25
CCU3	Jul-10	12.18	Calcium Carbonate	%CCE	AOAC 955.01	NA	NA
CCU3	Jul-10	15.86	Soluble Salts	$\begin{array}{ll} mmhos \ cm^{-1} & or \\ dS \ m^{-1} \end{array}$	Saturated paste extract	NA	4.0 after leaching
CCU3	Jul-10	7.3	Total Organic Carbon	% or g kg ⁻¹		NA	NA
CCU3	Jul-10	72.8	Particle Size Analysis	%Sand		NA	NA
CCU3	Jul-10	12.1	(<2 mm)	% Silt	<2 mm samples	NA	NA
CCU3	Jul-10	7.1	()	% Clay		NA	NA
CCU3	40381	8.0	Coarse fragments (>2 mm)		>2 mm samples	NA	NA

TABLE 9. SHIRLEY PLANTATION SCREENING TABLE Calvert Cliffs Sediment Characterization (2010)

NA= Indicates that criteria are not available.

1. For samples <RL, use 50% of RL for data entry column. One-half the RL is assumed for chemicals reported as non-detect or < RL; however, these values will not be used for exclusionary purposes unless other evidence indicates such. Values in *italics* are not "real" values, but an arbitrary entry.

2. Use **bold highlight** for all individual samples entered in working area and average sample values that exceed the "proposed VA upland fill criteria" in far right column. Highlight all values exceeding proposed VA exclusion criteria in **bold highlight red**. Put arbitrary values calculated as 50% the RL in *italics*. Tip: when copying numbers from your lab analytical results spreadsheets to this spreadsheet, samples with a "<" in front of them are typically at the RL and should reported as 50% RL and put in *italics*.

New Jersey Department of Environmental Protection, The Management and Regulation of Dredging Activities and Dredged Material in New Jersey's Tidal Waters. 1997. http://www.njstatelib.org/digit/r588/r5881997.html
 EPA Region 3 SSLs have been merged into a regional document developed with input from Regions III, VI, and IX. Values from September 12, 2008 version. Values listed for: antimony (metallic), arsenic (inorganic), chromium VI (particulates), lead and compounds, manganese and cadmium values are for diet, methyl mercury, nickel refinery dust, vanadium and compounds. Website: http://www.epa.gov/reg3hwmd/risk/human/rb-concentration_table/Gnerrie_Tables/index.htm

5. Background metal levels specific to the state of Virginia based on Smith, D.B. et al. 2005. Major- and Trace-Element Concentrations in Soils from Two Continental-Scale Transects of the United States and Canada. USGS Open File Report 2005-1253.http://pubs.usgs.gov/of/2005/1253/pdf/OFR1253.pdf

6. The proposed Virginia exclusion standards generally represent the higher of EPA RBC Industrial, NJDEP or EPA 503 EQ levels for a given parameter. Values exceeding these limits are questionable for acceptance. 7. Proposed VA clean fill criteria are based primarily on NJDEP residential cleanup criteria and manually adjusted for known issues with agricultural production/bioavailability. Values between the clean fill and exclusion criteria require a variation of the current management strategy.

8. Total Aroclor concentrations are reported as sum of seven individual aroclors.

9. Additional analyses for these basic properties are essential for determining the management or acceptance of dredge material.

Note: Minimum sampling is one composite sample per 50,000 yards of material in situ. A minimum of three samples per material is required regardless of volume. Specific information on sampling procedures should go into the brief descriptions box at the top of the spreadsheet.

TABLE 10. PHYSICAL CHARACTERISTICS OF SEDIMENTCALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

			CALVERT CLIFFS NUCLEAR POWER PLAN					
ANALYTE	UNITS	Average RL	CCU3-BAR-1/2- SED	CCU3-BAR-3/4- SED	CCU3-DIS- SED			
GRAVEL	%		2.1	2.4	19.6			
SAND	%		71.2	82.1	65			
SILT	%		18.6	9.3	8.5			
CLAY	%		8.1	6.2	6.9			
SILT+CLAY	%		26.7	15.5	15.4			
PERCENT SOLIDS	%	1	63.8	79.2	79.3			

There are no sediment quality guidelines for the physical characteristics parameters

NOTES:

RL = average reporting limit

TABLE 11. GENERAL CHEMISTRY CONCENTRATIONS IN SEDIMENTCALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

				CALVERT CLIFFS NUCLEAR POWER PLAN		
ANALYTE	UNITS	VCP Standard	Average RL	CCU3-BAR-1/2- SED	CCU3-BAR-3/4- SED	CCU3-DIS- SED
AMMONIA AS NITROGEN	MG/KG		6.8	137	24	16.6
NITRATE-NITRITE	MG/KG		1.4	0.76 B J	5.5 J	0.61 B J
РН				7.9	8.2	8.8
TOTAL CYANIDE	MG/KG		0.68	0.78 U	0.63 U	0.63 U
TOTAL KJELDAHL NITROGEN	MG/KG		204	749 J	426 J	319
TOTAL ORGANIC CARBON	%		0.13	1.51	0.17	0.52
TOTAL PHOSPHORUS	MG/KG		94	841	481	215
TOTAL SULFIDE	MG/KG		41	354	39.4	41.4
TOTAL PETROLEUM HYDROCARBON - DRO	UG/KG	230,000	140	370	40 J	130 U
TOTAL PETROLEUM HYDROCARBON - GRO	UG/KG	230,000	140	160 U	130 U	130 U

There are no sediment quality guidelines for the general chemistry parameters

NOTES: Bold values represent detected concentrations.

RL is reported for non-detected constituents.

RL = average reporting limit

B (inorganic) = compound was detected, but below the reporting limit (value is estimated)

 \mathbf{J} (inorganic) = detected in the laboratory method blank

J (organic) = compound was detected, but below the reporting limit (value is estimated)

TABLE 12. METAL CONCENTRATIONS (MG/KG) IN SEDIMENTCALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

				CALVERT CLIFFS NUCLEAR POWER PLANT				
ANALYTE	UNITS	Average MDL	ERL*	ERM*	CCU3-BAR-1/2- SED	CCU3-BAR-3/4- SED	CCU3-DIS- SED	
ALUMINUM	MG/KG	0.193			3,850	~	909	
						1,310		
ANTIMONY	MG/KG	0.002			0.084 B J	0.047 B J	0.067 B J	
ARSENIC	MG/KG	0.012	8.2	70.0	3.7	0.95	2.1	
BARIUM	MG/KG	0.007			16	6.6	15.8	
BERYLLIUM	MG/KG	0.005			0.26	0.11	0.065	
CADMIUM	MG/KG	0.005	1.2	9.6	0.38	0.22	0.45	
CALCIUM	MG/KG	0.893			31,100	30,600	89,300	
CHROMIUM	MG/KG	0.004	81	370	12.1 J	10.2 J	9.4 J	
COBALT	MG/KG	0.001			2.4	0.72	0.39	
COPPER	MG/KG	0.023	34	270	5.6	1.5	1.4	
IRON	MG/KG	0.240			8,140 J	2,570 J	1,950 J	
LEAD	MG/KG	0.003	47	218	7.4	2.4	1.3	
MAGNESIUM	MG/KG	0.130			3,050	1,170	1,140	
MANGANESE	MG/KG	0.007			143 J	44.5 J	13.6 J	
MERCURY	MG/KG	0.007	0.150	0.710	0.046	0.0069 U	0.0069 U	
NICKEL	MG/KG	0.008	20.9	51.6	9.6	3.7	3.4	
POTASSIUM	MG/KG	0.940			961	471	359	
SELENIUM	MG/KG	0.034			0.45	0.21 B	0.42	
SILVER	MG/KG	0.003	1.00	3.70	0.046 B	0.02 B	0.014 B	
SODIUM	MG/KG	0.940			2,760	1,410	1,840	
THALLIUM	MG/KG	0.001			0.15	0.13	0.2	
TIN	MG/KG	0.040			0.91 J	0.88 J	0.59 J	
VANADIUM	MG/KG	0.005			11.2 J	3.9 J	7.2 J	
ZINC	MG/KG	0.044	150	410	29.4	16.6	7.2	

*Source: MacDonald et al. 1996. Ecotoxicology 5: 253-278.

NOTES: Bold values represent detected concentrations.

MDL is reported for non-detected constituents.

MDL = average method detection limit

B (inorganic) = compound was detected, but below the reporting limit (value is estimated)

 \mathbf{J} (inorganic) = detected in the laboratory method blank

TABLE 13. PAH CONCENTRATIONS (UG/KG) IN SEDIMENTCALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

					CALVERT CLIFFS NUCLEAR POWER PLANT			
ANALYTE	UNITS	Average MDL	ERL*	ERM*	CCU3-BAR-1/2- SED	CCU3-BAR-3/4- SED	CCU3-DIS- SED	
2-METHYLNAPHTHALENE	UG/KG	2.07	70.00	670	9.2 J	1.9 U	1.9 U	
ACENAPHTHENE	UG/KG	2.17	16.00	500	14 J	2 U	2 U	
ACENAPHTHYLENE	UG/KG	2.60	44.00	640	3.5 J	2.4 U	2.4 U	
ANTHRACENE	UG/KG	2.23	85.30	1,100	8.9 J	2 U	2.1 U	
BENZO(A)ANTHRACENE	UG/KG	2.83	261	1,600	14 J	2.6 U	2.6 U	
BENZO(A)PYRENE	UG/KG	2.27	430	1,600	2.6 U	2.1 U	2.1 U	
BENZO(B)FLUORANTHENE	UG/KG	3.57			16 J	3.3 U	3.3 U	
BENZO(GHI)PERYLENE	UG/KG	2.27			2.6 U	2.1 U	2.1 U	
BENZO(K)FLUORANTHENE	UG/KG	4.60			9.8 J	4.2 U	4.3 U	
CHRYSENE	UG/KG	2.70	384	2,800	15 J	2.5 U	2.5 U	
DIBENZ(A,H)ANTHRACENE	UG/KG	2.50	63.4	260	2.9 U	2.3 U	2.3 U	
FLUORANTHENE	UG/KG	2.40	600	5,100	43	2.2 U	2.2 U	
FLUORENE	UG/KG	3.00	19	540	16 J	2.8 U	2.8 U	
INDENO(1,2,3-CD)PYRENE	UG/KG	2.37			2.7 U	2.2 U	2.2 U	
NAPHTHALENE	UG/KG	1.97	160	2,100	2.3 U	1.8 U	1.8 U	
PHENANTHRENE	UG/KG	3.60	240	1,500	44	3.3 U	3.3 U	
PYRENE	UG/KG	2.27	665	2,600	37	2.1 U	2.1 U	
TOTAL PAHs (ND=MDL)	UG/KG		4,022	44,792	237	21	21	

*Source: MacDonald et al. 1996. Ecotoxicology 5: 253-278.

NOTES: Bold values represent detected concentrations. Shaded concentrations exceed sediment quality guidelines.

MDL is reported for non-detected constituents.

MDL = average method detection limit

 \mathbf{J} (organic) = compound was detected, but below the reporting limit (value is estimated)

TABLE 14. PCB CONGENER CONCENTRATIONS (UG/KG) IN SEDIMENTCALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

					CALVERT CLIFFS NUCLEAR POWER PLANT			
		Average			CCU3-BAR-1/2-	CCU3-BAR-3/4-	CCU3-DIS-	
ANALYTE	UNITS	MDL	ERL**	ERM**	SED	SED	SED	
PCB 8 (BZ)	UG/KG	0.04			0.99 PG	0.044 U	0.082 J PG	
PCB 18 (BZ)	UG/KG	0.03			0.036 U	0.029 U	0.029 U	
PCB 28 (BZ)	UG/KG	0.05			0.38 PG	0.047 U	0.047 U	
PCB 44 (BZ)	UG/KG	0.04			0.73	0.043 U	0.043 U	
PCB 49 (BZ)	UG/KG	0.05			0.87	0.045 U	0.045 U	
PCB 52 (BZ)	UG/KG	0.04			1.4	0.042 U	0.042 U	
PCB 66 (BZ)	UG/KG	0.04			0.043 U	0.035 U	0.034 U	
PCB 77 (BZ)	UG/KG	0.05			0.057 U	0.046 U	0.046 U	
PCB 87 (BZ)	UG/KG	0.04			1.4 PG	0.039 U	0.039 U	
PCB 90 (BZ)	UG/KG	0.03			0.04 U	0.032 U	0.032 U	
PCB 101 (BZ)	UG/KG	0.04			2.3	0.043 U	0.043 U	
PCB 105 (BZ)	UG/KG	0.05			0.055 U	0.044 U	0.044 U	
PCB 118 (BZ)	UG/KG	0.04			1.6 PG	0.043 U	0.043 U	
PCB 126 (BZ)	UG/KG	0.06			0.069 U	0.055 U	0.055 U	
PCB 128 (BZ)	UG/KG	0.04			0.86	0.043 U	0.043 U	
PCB 138 (BZ)	UG/KG	0.05			3	0.045 U	0.045 U	
PCB 153 (BZ)	UG/KG	0.04			2.2	0.044 U	0.044 U	
PCB 156 (BZ)	UG/KG	0.04			0.45	0.043 U	0.043 U	
PCB 169 (BZ)	UG/KG	0.05			0.052 U	0.042 U	0.042 U	
PCB 170 (BZ)	UG/KG	0.04			0.43	0.043 U	0.043 U	
PCB 180 (BZ)	UG/KG	0.05			0.054 U	0.043 U	0.043 U	
PCB 183 (BZ)	UG/KG	0.04			0.19 J PG	0.042 U	0.042 U	
PCB 184 (BZ)	UG/KG	0.04			0.045 U	0.036 U	0.036 U	
PCB 187 (BZ)	UG/KG	0.05			0.28	0.045 U	0.045 U	
PCB 195 (BZ)	UG/KG	0.05			0.053 U	0.043 U	0.043 U	
PCB 206 (BZ)	UG/KG	0.05			0.052 U	0.042 U	0.042 U	
PCB 209 (BZ)	UG/KG	0.05			0.11 J	0.045 U	0.045 U	
TOTAL PCBS (ND=1/2MDL)	UG/KG		22.7	180	28.38	0.08	0.24	
TOTAL PCBS (ND=MDL)	UG/KG		22.7	180	29.07	1.55	1.63	

* PCB congeners used for Total PCB summation, as per Table 9-3 of the ITM (USEPA/USACE 1998)

**Source : MacDonald et al. 1996. Ecotoxicology 5: 253-278.

NOTES: Bold values represent detected concentrations. Shaded concentrations exceed sediment quality guidelines. MDL is reported for non-detected constituents.

MDL = average method detection limit

TABLE 15. PCB AROCLOR CONCENTRATIONS (UG/KG) IN SEDIMENT

CALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

			CALVERT CLIFFS NUCLEAR POWER PLANT					
ANALYTE	UNITS	Average MDL	CCU3-BAR-1/2- SED	CCU3-BAR-3/4- SED	CCU3-DIS- SED			
AROCLOR 1016	UG/KG	1.7	1.9 U	1.5 U	1.6 U			
AROCLOR 1221	UG/KG	2.2	2.5 U	2 U	2 U			
AROCLOR 1232	UG/KG	1.9	2.2 U	1.8 U	1.8 U			
AROCLOR 1242	UG/KG	1.8	2.1 U	1.7 U	1.7 U			
AROCLOR 1248	UG/KG	1.1	1.2 U	0.98 U	0.99 U			
AROCLOR 1254	UG/KG	1.6	22	1.5 U	1.5 U			
AROCLOR 1260	UG/KG	1.6	1.9 U	1.5 U	1.5 U			

There are no sediment quality guidelines for PCB Aroclors

NOTES: MDL is reported for non-detected constituents.

MDL = average method detection limit

TABLE 16. CHLORINATED PESTICIDE CONCENTRATIONS (UG/KG) IN SEDIMENTCALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

					CALVERT CLIFFS NUCLEAR POWER PLANT				
		Average	DDI 4			CCU3-BAR-3/4-	CCU3-DIS-		
ANALYTE	UNITS	MDL	ERL*	ERM*	SED	SED	SED		
4,4'-DDD	UG/KG	0.15	2	20	0.17 U	0.14 U	0.14 U		
4,4'-DDE	UG/KG	0.17	2.2	27	0.2 U	0.16 U	0.16 U		
4,4'-DDT	UG/KG	0.17	1	7	0.2 U	0.15 U	0.16 U		
ALDRIN	UG/KG	0.20			0.23 U	0.18 U	0.19 U		
ALPHA-BHC	UG/KG	0.18			0.29 J PG	0.17 U	0.17 U		
ALPHA-CHLORDANE	UG/KG	0.22	63.4	260	0.26 U	0.2 U	0.21 U		
BETA-BHC	UG/KG	0.29			0.34 U	0.27 U	0.27 U		
CHLORDANE (TECHNICAL)	UG/KG	0.50			0.58 U	0.45 U	0.46 U		
CHLOROBENSIDE	UG/KG	0.59			0.68 U	0.54 U	0.55 U		
DCPA	UG/KG	0.30			0.35 U	0.28 U	0.28 U		
DELTA-BHC	UG/KG	0.17			0.2 U	0.16 U	0.16 U		
DIELDRIN	UG/KG	0.19	0.02	8	0.22 U	0.17 U	0.18 U		
ENDOSULFAN I	UG/KG	0.21			0.25 U	0.19 U	0.2 U		
ENDOSULFAN II	UG/KG	0.20			0.23 U	0.49 J	0.48 J		
ENDOSULFAN SULFATE	UG/KG	0.12			0.14 U	0.11 U	0.11 U		
ENDRIN	UG/KG	0.22			1.1 J	0.2 U	0.2 U		
ENDRIN ALDEHYDE	UG/KG	0.22			0.25 U	0.2 U	0.2 U		
ENDRIN KETONE	UG/KG	0.17			0.2 U	0.16 U	0.16 U		
GAMMA-BHC (LINDANE)	UG/KG	0.20			0.41 J PG	0.38 J PG	0.45 J		
GAMMA-CHLORDANE	UG/KG	0.22			0.26 U	0.2 U	0.21 U		
HEPTACHLOR	UG/KG	0.25			0.37 J PG	0.23 U	0.54 J PG		
HEPTACHLOR EPOXIDE	UG/KG	0.22			0.25 U	0.2 U	0.2 U		
METHOXYCHLOR	UG/KG	0.24			0.27 U	0.22 U	0.22 U		
MIREX	UG/KG	0.10			0.12 U	0.095 U	0.097 U		
TOXAPHENE	UG/KG	7.53			8.7 U	6.9 U	7 U		

*Source: MacDonald et al. 1996. Ecotoxicology 5: 253-278.

NOTES: Bold values represent detected concentrations. Shaded concentrations exceed sediment quality guidelines.

MDL is reported for non-detected constituents.

MDL = average method detection limit

J (organic) = compound was detected, but below the reporting limit (value is estimated)

PG = the percent difference between the original and confirmation analysis is greater than 40%

TABLE 17. SVOC CONCENTRATIONS (UG/KG) IN SEDIMENTCALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

			CALVERT CLI	FFS NUCLEAR P	OWER PLANT
		Average	CCU3-BAR-1/2-	CCU3-BAR-3/4-	CCU3-DIS-
ANALYTE	UNITS	MDL	SED	SED	SED
1,2,4-TRICHLOROBENZENE	UG/KG	6.3	7.2 U	5.8 U	5.8 U
1,2-DICHLOROBENZENE	UG/KG	12.0	14 U	11 U	11 U
1,3-DICHLOROBENZENE	UG/KG	8.8	10 U	8.1 U	8.2 U
1,4-DICHLOROBENZENE	UG/KG	8.1	9.4 U	7.5 U	7.5 U
2,4,5-TRICHLOROPHENOL	UG/KG	12.0	14 U	11 U	11 U
2,4,6-TRICHLOROPHENOL	UG/KG	17.3	20 U	16 U	16 U
2,4-DICHLOROPHENOL	UG/KG	2.3	2.6 U	2.1 U	2.1 U
2,4-DIMETHYLPHENOL	UG/KG	17.3	20 U	16 U	16 U
2,4-DINITROPHENOL	UG/KG	136.7	160 U	120 U	130 U
2,4-DINITROTOLUENE	UG/KG	9.3	11 U	8.4 U	8.5 U
2,6-DINITROTOLUENE	UG/KG	11.7	13 U	11 U	11 U
2-CHLORONAPHTHALENE	UG/KG	2.4	2.7 U	2.2 U	2.2 U
2-CHLOROPHENOL	UG/KG	9.4	11 U	8.5 U	8.6 U
2-METHYLPHENOL	UG/KG	7.9	9.1 U	7.3 U	7.4 U
2-NITROANILINE	UG/KG	51.0	59 U	47 U	47 U
2-NITROPHENOL	UG/KG	12.7	14 U	12 U	12 U
3,3'-DICHLOROBENZIDINE	UG/KG	12.0	14 U	11 U	11 U
3-NITROANILINE	UG/KG	46.7	54 U	43 U	43 U
4,6-DINITRO-2-METHYLPHENOL	UG/KG	45.7	53 U	42 U	42 U
4-BROMOPHENYL PHENYL ETHER	UG/KG	9.8	11 U	9.1 U	9.2 U
4-CHLORO-3-METHYLPHENOL	UG/KG	10.4	12 U	9.6 U	9.7 U
4-CHLOROANILINE	UG/KG	8.9	10 U	8.4 U	8.4 U
4-CHLOROPHENYL PHENYL ETHER	UG/KG	13.0	15 U	12 U	12 U
4-METHYLPHENOL	UG/KG	11.0	13 U	10 U	10 U
4-NITROANILINE	UG/KG	46.0	53 U	42 U	43 U
4-NITROPHENOL	UG/KG	39.0	45 U	36 U	36 U
BIS(2-CHLOROETHOXY)METHANE	UG/KG	7.5	8.6 U	6.9 U	6.9 U
BIS(2-CHLOROETHYL) ETHER	UG/KG	3.0	3.5 U	2.8 U	2.8 U
BIS(2-CHLOROISOPROPYL) ETHER	UG/KG	2.5	2.8 U	2.3 U	2.3 U
BIS(2-ETHYLHEXYL) PHTHALATE	UG/KG	18.3	43 J	17 U	32 J
BUTYL BENZYL PHTHALATE	UG/KG	15.3	18 U	14 U	14 U
CARBAZOLE	UG/KG	2.1	2.4 U	1.9 U	1.9 U
DIBENZOFURAN	UG/KG	11.0	13 U	10 U	10 U
DIETHYL PHTHALATE	UG/KG	12.0	14 U	11 U	11 U
DIMETHYL PHTHALATE	UG/KG	12.0	14 U	11 U	11 U
DI-N-BUTYL PHTHALATE	UG/KG	14.0	16 U	13 U	13 U
DI-N-OCTYL PHTHALATE	UG/KG	12.0	14 U	11 U	11 U
HEXACHLOROBENZENE	UG/KG	2.4	2.8 U	2.2 U	2.2 U
HEXACHLOROBUTADIENE	UG/KG	2.5	2.9 U	2.3 U	2.4 U
HEXACHLOROCYCLOPENTADIENE	UG/KG	12.0	14 U	11 U	11 U
HEXACHLOROETHANE	UG/KG	8.2	9.4 U	7.5 U	7.6 U
ISOPHORONE	UG/KG	8.6	9.9 U	7.9 U	7.9 U
NITROBENZENE	UG/KG	9.5	11 U	8.7 U	8.8 U
N-NITROSODI-N-PROPYLAMINE	UG/KG	2.7	3.1 U	2.5 U	2.5 U
N-NITROSODIPHENYLAMINE	UG/KG	10.5	12 U	9.7 U	9.7 U
PENTACHLOROPHENOL	UG/KG	10.2	12 U	9.3 U	9.4 U
PHENOL	UG/KG	2.7	79	5.9 J	2.5 U

There are no sediment quality guidelines for the general chemistry parameters

NOTES: Bold values represent detected concentrations.

MDL is reported for non-detected constituents.

MDL = average method detection limit

J (organic) = compound was detected, but below the reporting limit (value is estimated)

TABLE 18. BUTYLTIN AND 2,3,7,8-TCDD CONCENTRATIONS (UG/KG) IN
SEDIMENT

CALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

			CALVERT CLIFFS NUCLEAR POWER PLANT				
ANALYTE	UNITS	Average RL	CCU3-BAR-1/2- SED	CCU3-BAR-3/4- SED	CCU3-DIS- SED		
DIBUTYLTIN	UG/KG	1.7	1.9 U	1.7 U	1.6 U		
MONOBUTYLTIN	UG/KG	6.7	7.3 U	6.4 U	6.3 U		
TETRABUTYLTIN	UG/KG	2.3	2.5 U	2.2 U	2.1 U		
TRIBUTYLTIN	UG/KG	2.0	2.2 U	1.9 U	1.9 U		
2,3,7,8-TCDD	PG/G	0.25	0.2 U	0.23 U	0.33 U		

NOTES: RL is reported for non-detected constituents.

RL = average reporting limit

TABLE 19. GENERAL CHEMISTRY CONCENTRATIONS IN SITE WATER AND STANDARD AND EFFLUENT ELUTRIATES CALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

ANALYTE	UNITS	Average RL	USEPA ACUTE CRITERIA	USEPA CHRONIC CRITERIA	Site Water
AMMONIA AS NITROGEN	MG/L	0.23	3	0.45	3.7
DISSOLVED CYANIDE	UG/L	10			10 U
DISSOLVED NITRATE/NITRITE	MG/L	0.10			
TOTAL NITRATE/NITRITE	MG/L	1.00			0.025 B J
DISSOLVED ORGANIC CARBON	MG/L	1			2.7
DISSOLVED SULFIDE	MG/L	3			
TOTAL SULFIDE	MG/L	3			3 U
TOTAL KJELDAHL NITROGEN	MG/L	3			10.7
PHOSPHORUS AS ORTHOPHOSPHATE	MG/L	0.1			
TOTAL PHOSPHORUS	MG/L	0.10			0.1 U
TPH (AS DIESEL)	UG/L	100			200 B
TPH (AS GASOLINE)	UG/L	100			100 U

St	Standard Elutriate									
CCU3-BAR- 1/2-SET	CCU3-BAR- 3/4-SET	CCU3-DIS- SET								
26.4	5.5	4.9								
10 U	10 U	10 U								
0.053 B J	3.1 J	1.9 J								
3.1	1.5	1.6								
3 U	3 U	3 U								
29.2 J	7.3 J	3.4 J								
0.1 U	0.039 B	0.1 U								
230 B	55 J B	73 J B								
100 U	100 U	100 U								

	E	Effluent Elutriate									
ois-	CCU3-BAR- 1/2-EFF	CCU3-BAR- 3/4-EFF	CCU3-DIS- EFF								
	17.4	3.1	2.9								
	10 U	10 U	10 U								
	0.029 B J	1.8 J	0.22 J								
	2.4	1.2	1.4								
	3 U	3 U	3 U								
	15.7 J	5.6 J	2.8 B J								
T											
	0.1 U	0.1 U	0.1 U								
3	61 J	78 J	54 J								
J	100 U	100 U	100 U								

Source : USEPA 2010. National Recommended Water Quality Criteria

NOTES: Bold values represent detected concentrations, shaded values exceed acute or chronic criteria.

RL is reported for non-detected constituents.

RL = average reporting limit

 \mathbf{B} (inorganic) = compound was detected, but below the reporting limit (value is estimated)

 \mathbf{J} (inorganic) = detected in the laboratory method blank

B (organic) = detected in the laboratory method blank

J (organic) = compound was detected, but below the reporting limit (value is estimated)

TABLE 20. METAL CONCENTRATIONS (UG/L) IN SITE WATER AND STANDARD AND EFFLUENT ELUTRIATES	
CALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)	

						Standard Elutriate		Ef	Effluent Elutriate		
ANALYTE	UNITS	Average MDL	USEPA ACUTE CRITERIA	USEPA CHRONIC CRITERIA	Site Water	CCU3-BAR- 1/2-SET	CCU3-BAR- 3/4-SET	CCU3-DIS- SET	CCU3- BAR-1/2- EFF	CCU3- BAR-3/4- EFF	CCU3-DIS- EFF
ALUMINUM	UG/L	12.8			2.6 U	12.8 U	12.8 U	12.8 U	22 B	12.8 U	12.8 U
ANTIMONY	UG/L	0.094			0.1 B	3 B	0.92 B	1 B	1.1 B	0.9 B	0.7 B
ARSENIC	UG/L	1.5	69	36	4.1	5.3	5	6.5	6.2	7.2	8.5
BARIUM	UG/L	0.49			30.1	187	37.4 B	36.2 B	142	34.7 B	33.2 B
BERYLLIUM	UG/L	0.18			0.037 U	0.18 U	0.18 U	0.18 U	0.18 U	0.18 U	0.18 U
CADMIUM	UG/L	0.57	40	8.8	0.11 U	0.57 U	0.57 U	0.57 U	0.57 U	0.57 U	0.57 U
CALCIUM	UG/L	14.2			178,000 J	154,000	198,000	180,000	158,000	176,000	174,000
CHROMIUM	UG/L	2.7	1100	50	5.8	7.3 B	7.7 B	7.8 B	8.9 B J	7.5 B J	7.4 B J
COBALT	UG/L	0.13			0.83	0.88 B	0.8 B	0.83 B	0.74 B	0.62 B	0.84 B
COPPER	UG/L	1.2	4.8	3.1	2.6	2.7 B	9.7 B	3.7 B	3.2 B J	5.2 B J	4.5 B J
IRON	UG/L	30.5			87.5	116 B	67.9 B	60.5 B	70.4 B	77 B	63.3 B
LEAD	UG/L	0.096	210	8.1	0.44 B	0.096 U	0.2 B	0.096 U	0.096 U	0.096 U	0.24 B
MAGNESIUM	UG/L	5.8			480,000	495,000	506,000	493,000	492,000 J	486,000 J	485,000 J
MANGANESE	UG/L	0.19			50.3	52.9	4.2	9.5	18.2	1.9 B	5.7
MERCURY	UG/L	0.038	1.8	0.94	0.038 U	0.038 U	0.038 U	0.038 U	0.038 U	0.038 U	0.038 U
NICKEL	UG/L	0.87	74	8.2	1.3	5.5	6.4	9.7	4.9 B	4.3 B	7.1
POTASSIUM	UG/L	29.1			152,000	160,000 J	164,000 J	157,000 J	161,000 J	156,000 J	154,000 J
SELENIUM	UG/L	2.1	290	71	19.8	20.9 B	22.9 B	27.4	25	25.3	34.3
SILVER	UG/L	0.18	1.9		0.036 U	0.18 U	0.18 U	0.18 U	0.18 U	0.18 U	0.18 U
SODIUM	UG/L	191			4,250,000	3,980,000 J	3,920,000 J	4,000,000 J	3,920,000 J	3,830,000 J	3,840,000 J
THALLIUM	UG/L	0.076			0.042 B J	0.14 B	0.09 B	0.076 U	0.74 B J	0.076 U	0.08 B J
TIN	UG/L	7.5			1.5 U	7.5 U	7.5 U	7.5 U	9.6 B	7.5 U	7.5 U
VANADIUM	UG/L	0.41			1.9 J	1.4 B	0.41 U	0.41 U	0.41 U	1.6 B	1.2 B
ZINC	UG/L	4.8	90	81	6	7 B	14.6 B	9.3 B	6.8 B	15.6 B	6.5 B

Source : USEPA 2010. National Recommended Water Quality Criteria

NOTES: Bold values represent detected concentrations, shaded values exceed acute or chronic criteria.

MDL is reported for non-detected constituents.

MDL = average method detection limit

B (inorganic) = compound was detected, but below the reporting limit (value is estimated)

 \mathbf{J} (inorganic) = detected in the laboratory method blank

TABLE 21. PAH CONCENTRATIONS (UG/L) IN SITE WATER AND STANDARD AND EFFLUENT ELUTRIATES CALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

ANALYTE	UNITS	Average MDL	Site Water
2-METHYLNAPHTHALENE	UG/L	0.011	0.023 J B
ACENAPHTHENE	UG/L	0.014	0.031 J
ACENAPHTHYLENE	UG/L	0.014	0.014 U
ANTHRACENE	UG/L	0.014	0.014 U
BENZO(A)ANTHRACENE	UG/L	0.014	0.014 U
BENZO(A)PYRENE	UG/L	0.013	0.013 U
BENZO(B)FLUORANTHENE	UG/L	0.015	0.015 J
BENZO(GHI)PERYLENE	UG/L	0.014	0.014 U
BENZO(K)FLUORANTHENE	UG/L	0.051	0.051 U
CHRYSENE	UG/L	0.013	0.013 U
DIBENZ(A,H)ANTHRACENE	UG/L	0.015	0.015 U
FLUORANTHENE	UG/L	0.015	0.015 U
FLUORENE	UG/L	0.020	0.032 J B
INDENO(1,2,3-CD)PYRENE	UG/L	0.019	0.019 U
NAPHTHALENE	UG/L	0.013	0.036 J
PHENANTHRENE	UG/L	0.040	0.089 J B
PYRENE	UG/L	0.015	0.015 U
TOTAL PAHS (ND=MDL)	UG/L		0.325

Standard Elutriate							
CCU3-BAR- 1/2-SET	CCU3-BAR- 3/4-SET	CCU3-DIS- SET					
0.014 J	0.011 U	0.011 U					
0.014 U	0.014 U	0.014 U					
0.014 U	0.014 U	0.014 U					
0.014 U	0.014 U	0.014 U					
0.014 U	0.014 U	0.014 U					
0.013 U	0.013 U	0.013 U					
0.015 U	0.015 U	0.015 U					
0.014 U	0.014 U	0.014 U					
0.051 U	0.051 U	0.051 U					
0.013 U	0.013 U	0.013 U					
0.015 U	0.015 U	0.015 U					
0.015 U	0.015 U	0.015 U					
0.02 U	0.02 U	0.02 U					
0.019 U	0.019 U	0.019 U					
0.018 J	0.013 U	0.013 U					
0.061 J	0.042 J	0.044 J					
0.015 U	0.015 U	0.015 U					
0.216	0.177	0.179					

1	Effluent Elutriate								
1	CCU3-		CCU3-DIS-						
	BAR-1/2-	BAR-3/4-	EFF						
	0.013 J	0.011 U	0.012 U						
	0.048 J	0.014 U	0.014 U						
	0.014 U	0.014 U	0.014 U						
	0.014 U	0.014 U	0.015 U						
	0.18 J	0.014 U	0.014 U						
	0.12 J	0.013 U	0.013 U						
	0.16 J	0.015 U	0.015 U						
	0.45	0.014 U	0.014 U						
	0.23	0.051 U	0.052 U						
	0.26	0.013 U	0.013 U						
	0.52	0.015 U	0.015 U						
	0.015 U	0.015 U	0.015 U						
	0.027 J	0.02 U	0.021 U						
	0.53	0.019 U	0.019 U						
	0.013 U	0.019 J	0.013 U						
	0.058 J	0.054 J	0.041 U						
	0.023 J	0.015 U	0.015 U						
1	2.647	0.202	0.158						

NOTES: There are no USEPA acute or chronic water quality criteria for PAHs

Bold values represent detected concentrations.

MDL is reported for non-detected constituents

MDL = average method detection limit

J (organic) = compound was detected, but below the reporting limit (value is estimated)

					Standard Elutriate			Effluent Elutriate		
ANALYTE	UNITS	Average MDL	USEPA CHRONIC CRITERIA	Site Water	CCU3-BAR- 1/2-SET	CCU3-BAR- 3/4-SET	CCU3-DIS- SET	CCU3- BAR-1/2- EFF	CCU3- BAR-3/4- EFF	CCU3-DIS- EFF
PCB 8 (BZ) (a) (b)	NG/L	0.36		0.41 U	0.36 U	0.36 U	0.36 U	0.36 U	0.36 U	0.36 U
PCB 18 (BZ) (a) (b)	NG/L	0.36		0.45 U	0.58 J PG	0.92 J PG	0.36 U	0.36 U	0.36 U	0.36 U
PCB 28 (BZ) (a) (b)	NG/L	0.42		0.41 U	0.42 U	0.42 U	0.42 U	0.42 U	0.42 U	0.42 U
PCB 44 (BZ) ^{(a) (b)}	NG/L	0.43		0.41 U	1.6 PG	0.43 U	0.43 U	0.43 U	0.43 U	0.43 U
PCB 49 (BZ) (a)	NG/L	0.26		0.42 U	0.26 U	0.26 U	0.26 U	0.26 U	0.26 U	0.26 U
PCB 52 (BZ) ^{(a) (b)}	NG/L	0.41		0.4 U	0.41 U	0.41 U	0.41 U	0.41 U	0.41 U	0.41 U
PCB 66 (BZ) ^{(a) (b)}	NG/L	0.46		0.47 U	0.46 U	0.46 U	0.46 U	0.46 U	0.46 U	0.46 U
PCB 77 (BZ) (a)	NG/L	0.45		0.41 U	8	0.45 U	0.45 U	0.45 U	0.45 U	0.45 U
PCB 87 (BZ) (a)	NG/L	0.41		0.38 U	0.41 U	0.41 U	0.41 U	0.41 U	0.41 U	0.41 U
PCB 90 (BZ)	NG/L	0.42		0.73 U	0.42 U	0.42 U	0.42 U	0.42 U	0.42 U	0.42 U
PCB 101 (BZ) (a) (b)	NG/L	0.45		0.39 U	0.45 U	0.45 U	0.45 U	0.45 U	0.45 U	0.45 U
PCB 105 (BZ) ^{(a) (b)}	NG/L	0.44		0.36 U	0.44 U	0.44 U	0.44 U	0.44 U	0.44 U	0.44 U
PCB 118 (BZ) (a) (b)	NG/L	0.46		0.5 U	0.46 U	0.46 U	0.46 U	0.46 U	0.46 U	0.46 U
PCB 126 (BZ) ^(a)	NG/L	0.3		0.37 U	0.3 U	0.3 U	0.3 U	0.3 U	0.3 U	0.3 U
PCB 128 (BZ) ^{(a) (b)}	NG/L	0.47		0.33 U	0.47 U	0.47 U	0.47 U	0.47 U	0.47 U	0.47 U
PCB 138 (BZ) (a) (b)	NG/L	0.46		0.32 U	0.46 U	0.46 U	0.46 U	0.72 J	0.46 U	0.46 U
PCB 153 (BZ) (a) (b)	NG/L	0.43		0.37 U	0.43 U	0.43 U	0.43 U	0.43 U	0.43 U	0.43 U
PCB 156 (BZ) ^(a)	NG/L	0.41		0.35 U	0.41 U	0.41 U	0.41 U	0.41 U	0.41 U	0.41 U
PCB 169 (BZ) ^(a)	NG/L	0.23		0.4 U	0.23 U	0.23 U	0.23 U	0.23 U	0.23 U	0.23 U
PCB 170 (BZ) (a) (b)	NG/L	0.22		0.35 U	0.22 U	0.22 U	0.22 U	0.22 U	0.22 U	0.22 U
PCB 180 (BZ) (a) (b)	NG/L	0.28		0.34 U	0.28 U	0.28 U	0.28 U	0.28 U	0.28 U	0.28 U
PCB 183 (BZ) ^(a)	NG/L	0.47		0.35 U	0.47 U	0.47 U	0.47 U	0.47 U	0.47 U	0.47 U
PCB 184 (BZ) ^(a)	NG/L	0.22		0.4 U	0.22 U	0.22 U	0.67 J PG	0.22 U	0.22 U	0.22 U
PCB 187 (BZ) (a) (b)	NG/L	0.46		0.37 U	0.46 U	0.46 U	0.46 U	0.46 U	0.46 U	0.46 U
PCB 195 (BZ) (a) (b)	NG/L	0.27		0.37 U	0.27 U	0.27 U	0.27 U	0.27 U	0.27 U	0.27 U
PCB 206 (BZ) ^{(a) (b)}	NG/L	0.29		0.36 U	0.29 U	0.29 U	0.29 U	0.29 U	0.29 U	0.29 U
PCB 209 (BZ) ^{(a) (b)}	NG/L	0.25		0.41 U	0.25 U	0.25 U	0.25 U	0.25 U	0.25 U	0.25 U
TOTAL PCBs (ND=MDL)	NG/L		30	14.1	14.7	14.2	14.2	14.7	14.2	14.2
*PCB congeners used for Total PCB sun	mation, as per T	able 9-3 of t	he ITM (USEPA	(USACE 1998)						

TABLE 22. PCB CONGENER* CONCENTRATIONS (NG/L) IN SITE WATER AND STANDARD AND EFFLUENT ELUTRIATES CALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

*PCB congeners used for Total PCB summation, as per Table 9-3 of the ITM (USEPA/USACE 1998)

Source : USEPA 2010. National Recommended Water Quality Criteria

NOTES: There are no USEPA acute water quality criteria for PCBs

Bold values represent detected concentrations, shaded values exceed chronic criteria.

MDL is reported for non-detected constituents

MDL = average method detection limit

TABLE 23. PCB AROCLOR CONCENTRATIONS (UG/L) IN SITE WATER AND STANDARD AND EFFLUENT ELUTRIATES CALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

ANALYTE	UNITS	Average MDL	Site Water
AROCLOR 1016	UG/L	0.10	0.095 U
AROCLOR 1221	UG/L	0.095	0.094 U
AROCLOR 1232	UG/L	0.11	0.11 U
AROCLOR 1242	UG/L	0.071	0.07 U
AROCLOR 1248	UG/L	0.086	0.085 U
AROCLOR 1254	UG/L	0.087	0.086 U
AROCLOR 1260	UG/L	0.051	0.051 U

Standard Elutriate						
CCU3-BAR-	CCU3-BAR-	CCU3-DIS-				
1/2-SET	3/4-SET	SET				
0.096 U	0.096 U	0.096 U				
0.095 U	0.095 U	0.095 U				
0.11 U	0.11 U	0.11 U				
0.071 U	0.071 U	0.071 U				
0.086 U	0.086 U	0.086 U				
0.087 U	0.087 U	0.087 U				
0.051 U	0.051 U	0.051 U				

Effluent Elutriate							
CCU3- CCU3- CCU3-DIS-							
BAR-1/2-	BAR-3/4-	EFF					
0.36 U	0.36 U	0.36 U					
0.36 U	0.36 U	0.36 U					
0.42 U	0.42 U	0.42 U					
0.43 U	0.43 U	0.43 U					
0.26 U	0.26 U	0.26 U					
0.41 U	0.41 U	0.41 U					
0.46 U	0.46 U	0.46 U					

NOTES: There are no USEPA acute or chronic water quality criteria for PCB Aroclors. Bold values represent detected concentrations.

MDL is reported for non-detected constituents

MDL = average method detection limit

TABLE 24. CHLORINATED PESTICIDE CONCENTRATIONS (UG/L) IN SITE WATER AND STANDARD AND EFFLUENT ELUTRIATES CALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

ANALYTE	UNITS	Average MDL	USEPA ACUTE CRITERIA	USEPA CHRONIC CRITERIA	Site Water
4,4'-DDD	UG/L	0.026			0.025 U
4,4'-DDE	UG/L	0.03			0.03 U
4,4'-DDT	UG/L	0.028	0.13	0.001	0.028 U
ALDRIN	UG/L	0.032	1.3		0.031 U
ALPHA-BHC	UG/L	0.025			0.025 U
ALPHA-CHLORDANE	UG/L	0.037			0.037 U
BETA-BHC	UG/L	0.038			0.037 U
CHLORDANE (TECHNICAL)	UG/L	0.063			0.062 U
CHLOROBENSIDE	UG/L	0.056			0.056 U
DCPA	UG/L	0.013			0.013 U
DELTA-BHC	UG/L	0.017			0.016 U
DIELDRIN	UG/L	0.031	0.71	0.0019	0.031 U
ENDOSULFAN I	UG/L	0.036	0.034	0.0087	0.035 U
ENDOSULFAN II	UG/L	0.037	0.034	0.0087	0.037 U
ENDOSULFAN SULFATE	UG/L	0.022			0.021 U
ENDRIN	UG/L	0.037	0.037	0.0023	0.036 U
ENDRIN ALDEHYDE	UG/L	0.034			0.034 U
ENDRIN KETONE	UG/L	0.035			0.035 U
GAMMA-BHC (LINDANE)	UG/L	0.03	0.16		0.03 U
GAMMA-CHLORDANE	UG/L	0.037			0.036 U
HEPTACHLOR	UG/L	0.038	0.053	0.0036	0.037 U
HEPTACHLOR EPOXIDE	UG/L	0.037	0.053	0.0036	0.037 U
METHOXYCHLOR	UG/L	0.035		0.03	0.034 U
MIREX	UG/L	0.018		0.001	0.018 U
TOXAPHENE	UG/L	0.71	0.21	0.0002	0.7 U

Si	tandard Elutria	ite
CCU3-BAR- 1/2-SET	CCU3-BAR- 3/4-SET	CCU3-DIS- SET
0.026 U	0.026 U	0.026 U
0.03 U	0.03 U	0.03 U
0.028 U	0.028 U	0.028 U
0.032 U	0.032 U	0.032 U
0.025 U	0.025 U	0.025 U
0.037 U	0.037 U	0.037 U
0.038 U	0.038 U	0.15
0.063 U	0.063 U	0.063 U
0.056 U	0.056 U	0.056 U
0.013 U	0.013 U	0.013 U
0.017 U	0.017 U	0.017 U
0.031 U	0.031 U	0.031 U
0.036 U	0.036 U	0.036 U
0.037 U	0.037 U	0.037 U
0.022 U	0.022 U	0.022 U
0.037 U	0.037 U	0.037 U
0.034 U	0.034 U	0.034 U
0.035 U	0.035 U	0.035 U
0.03 U	0.03 U	0.03 U
0.037 U	0.037 U	0.037 U
0.038 U	0.038 U	0.038 U
0.037 U	0.037 U	0.037 U
0.035 U	0.035 U	0.035 U
0.018 U	0.018 U	0.018 U
0.71 U	0.71 U	0.71 U

Efi	Effluent Elutriate						
CCU3-BAR- 1/2-EFF	CCU3- BAR-3/4- EFF	CCU3-DIS- EFF					
0.0003 U	0.0003 U	0.0003 U					
0.0013 U	0.0013 U	0.0013 U					
0.0003 U	0.0003 U	0.0003 U					
0.0015 U	0.0015 U	0.0015 U					
0.00045 J PG	0.00042 U	0.00042 U					
0.0014 U	0.0014 U	0.0014 U					
0.0016 U	0.0016 U	0.0016 U					
0.0018 J PG	0.0013 U	0.0013 U					
0.0067 PG	0.0048 PG	0.0063 PG					
0.0031 U	0.0031 U	0.0031 U					
0.0028 U	0.0028 U	0.0028 U					
0.00065 U	0.00065 U	0.00065 U					
0.00084 U	0.00084 U	0.00084 U					
0.0016 U	0.0016 U	0.0016 U					
0.0018 U	0.0018 U	0.0018 U					
0.0019 U	0.0019 U	0.0019 U					
0.0011 U	0.0011 U	0.0011 U					
0.0018 U	0.0018 U	0.0018 U					
0.0017 U	0.0017 U	0.0017 U					
0.0025	0.0024 J	0.0016 J PG					
0.0085	0.01	0.0083					
0.0018 U	0.0018 U	0.0018 U					
0.0017 U	0.0017 U	0.0017 U					
0.00091 U	0.00091 U	0.00091 U					
0.0013 U	0.0013 U	0.0013 U					

Source : USEPA 2010. National Recommended Water Quality Criteria

NOTES: Bold values represent detected concentrations, shaded values exceed acute or chronic criteria MDL is reported for non-detected constituents

MDL = average method detection limit

J (organic) = compound was detected, but below the reporting limit (value is estimated)

 \mathbf{PG} = the percent difference between the original and confirmation analysis is greater than 40%

TABLE 25. SVOC CONCENTRATIONS (UG/L) IN SITE WATER AND STANDARD AND EFFLUENT ELUTRIATES CALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

Average ANALYTE USEPA WDL USEPA CRITERIA USEPA CHRONIC CRITERIA Site Water CRITERIA CCU3-BAR- 3/4-SET CCU3-BAR- SET CCU3- SET BAR-1/2- SET BAR-1/2- SET BAR-3/4 EFF 1,2-DICHLOROBENZENE UG/L 0.067 0.067 U 0.067 U 0.067 U 0.067 U 0.067 U 0.067 U 0.07 U 1,3-DICHLOROBENZENE UG/L 0.07 0.07 U 0.07 U	CCU3-DIS- EFF 0.068 U 0.071 U 0.071 U 0.071 U 0.15 U 0.17 U 0.032 U 0.081 U 0.58 U 0.051 U 0.076 U 0.014 U 0.16 U
1.2-DICHLOROBENZENE UG/L 0.07 0.07 U 0.07 U 0.07 U 0.07 U 0.07 U 1,3-DICHLOROBENZENE UG/L 0.07 0.07 U 0.07 U <t< th=""><th>0.071 U 0.071 U 0.071 U 0.15 U 0.17 U 0.032 U 0.081 U 0.58 U 0.051 U 0.076 U 0.014 U</th></t<>	0.071 U 0.071 U 0.071 U 0.15 U 0.17 U 0.032 U 0.081 U 0.58 U 0.051 U 0.076 U 0.014 U
1,3-DICHLOROBENZENE UG/L 0.07 0.07 U 0.07 U 0.07 U 0.07 U 1,4-DICHLOROBENZENE UG/L 0.07 0.07 U 0.07 U 0.07 U 0.07 U 0.07 U 2,4,5-TRICHLOROPHENOL UG/L 0.14 0.14 U 0.03 U 0.03 U 0.03 U 0.03	0.071 U 0.071 U 0.15 U 0.17 U 0.032 U 0.081 U 0.58 U 0.051 U 0.076 U 0.014 U
1.3-DICHLOROBENZENE UG/L 0.07 0.07 U 0.07 U 0.07 U 0.07 U 0.07 U 1.4-DICHLOROBENZENE UG/L 0.07 0.07 U 0.01 U 0.03	0.071 U 0.071 U 0.15 U 0.17 U 0.032 U 0.081 U 0.58 U 0.051 U 0.076 U 0.014 U
1.4-DICHLOROBENZENE UG/L 0.07 0.07 U 0.03 U <th0< td=""><td>0.071 U 0.15 U 0.17 U 0.032 U 0.081 U 0.58 U 0.051 U 0.076 U 0.014 U</td></th0<>	0.071 U 0.15 U 0.17 U 0.032 U 0.081 U 0.58 U 0.051 U 0.076 U 0.014 U
2,4,5-TRICHLOROPHENOL UG/L 0.14 0.14 U 0.13 U </td <td>0.15 U 0.17 U 0.032 U 0.081 U 0.58 U 0.051 U 0.076 U 0.014 U</td>	0.15 U 0.17 U 0.032 U 0.081 U 0.58 U 0.051 U 0.076 U 0.014 U
2.4-DICHLOROPHENOL UG/L 0.031 0.031 U	0.17 U 0.032 U 0.081 U 0.58 U 0.051 U 0.076 U 0.014 U
2,4-DIMETHYLPHENOL UG/L 0.08 0.08 U 0.05 U 0.075 U	0.081 U 0.58 U 0.051 U 0.076 U 0.014 U
2,4-DINITROPHENOL UG/L 0.58 0.58 U 0.58 U 0.58 U 0.58 U 0.58 U 2,4-DINITROTOLUENE UG/L 0.05 0.05 U 0.075 U <td< td=""><td>0.58 U 0.051 U 0.076 U 0.014 U</td></td<>	0.58 U 0.051 U 0.076 U 0.014 U
2,4-DINITROTOLUENE UG/L 0.05 0.05 U 0.075	0.051 U 0.076 U 0.014 U
2,6-DINITROTOLUENE UG/L 0.075 0.075 U	0.076 U 0.014 U
	0.014 U
	0.16 U
2-CHLOROPHENOL UG/L 0.16 0.16 U 0.16 U 0.16 U 0.16 U 0.16 U 0.16 U	
2-METHYLPHENOL UG/L 0.081 0.081 U	0.082 U
2-NITROANILINE UG/L 0.33 0.33 U 0.33 U 0.33 U 0.33 U 0.33 U 0.33 U	0.33 U
2-NITROPHENOL UG/L 0.16 0.16 U 0.16 U 0.16 U 0.16 U 0.16 U 0.16 U	0.16 U
3,3'-DICHLOROBENZIDINE UG/L 0.11 0.11 U 0.11 U 0.11 U 0.11 U 0.11 U 0.11 U	0.11 U
3-NITROANILINE UG/L 0.3 0.3 U	0.31 U
4,6-DINITRO-2-METHYLPHENOL UG/L 0.21 0.21 U	0.21 U
4-BROMOPHENYL PHENYL ETHER UG/L 0.06 0.06 U 0.06 U 0.06 U 0.06 U 0.06 U 0.06 U	0.06 U
4-CHLORO-3-METHYLPHENOL UG/L 0.071 0.071 U	0.072 U
4-CHLOROANILINE UG/L 0.083 0.083 U	0.084 U
4-CHLOROPHENYL PHENYL ETHER UG/L 0.047 0.047 U	0.048 U
4-METHYLPHENOL UG/L 0.085 0.085 U 0.085 U 0.085 U 0.085 U 0.085 U 0.085 U	0.086 U
4-NITROANILINE UG/L 0.16 0.16 U 0.16 U 0.16 U 0.16 U 0.16 U 0.16 U	0.16 U
4-NITROPHENOL UG/L 0.57 0.57 U 0.57 U 0.57 U 0.57 U 0.57 U 0.57 U	0.57 U
BIS(2-CHLOROETHOXY)METHANE UG/L 0.055 0.055 U	0.055 U
BIS(2-CHLOROETHYL) ETHER UG/L 0.024 0.024 U	0.024 U
BIS(2-CHLOROISOPROPYL) ETHER UG/L 0.019 0.019 U	0.019 U
BIS(2-ETHYLHEXYL) PHTHALATE UG/L 0.75 1.5 J B 0.75 U 0.89 J 0.75 U 1.1 1.2	0.76 U
BUTYL BENZYL PHTHALATE UG/L 0.13 0.13 U	0.14 U
CARBAZOLE UG/L 0.015 0.015 U 0.015 U 0.015 U 0.015 U 0.015 U 0.015 U	0.015 U
DIBENZOFURAN UG/L 0.058 0.058 U 0.058 U 0.058 U 0.058 U 0.058 U	0.059 U
DIETHYL PHTHALATE UG/L 0.14 0.14 U 0.15 J 0.14 U 0.14 U 0.14 U 0.14 U 0.14 U	0.14 U
DIMETHYL PHTHALATE UG/L 0.072 0.072 U	0.073 U
DI-N-BUTYL PHTHALATE UG/L 0.12 0.12 U 0.12 U 0.12 U 0.12 U 0.12 U 0.12 U	0.12 U
DI-N-OCTYL PHTHALATE UG/L 0.19 0.19 U	0.2 U
HEXACHLOROBENZENE UG/L 0.017 0.017 U	0.017 U
HEXACHLOROBUTADIENE UG/L 0.016 0.016 U	0.016 U
HEXACHLOROCYCLOPENTADIENE UG/L 0.049 0.049 U	0.049 U
HEXACHLOROETHANE UG/L 0.059 0.059 U 0.059 U 0.059 U 0.059 U 0.059 U 0.059 U	0.06 U
ISOPHORONE UG/L 0.061 0.061 U	0.061 U
NITROBENZENE UG/L 0.079 0.079 U	0.08 U
N-NITROSODI-N-PROPYLAMINE UG/L 0.029 0.029 U	0.029 U
N-NITROSODIPHENYLAMINE UG/L 0.08 0.08 U	0.081 U
PENTACHLOROPHENOL UG/L 0.062 13 7.9 0.062 U 0.	0.063 U
PHENOL UG/L 0.0550 0.055 U 7.8 0.055 U	0.055 U

Source : USEPA 2010. National Recommended Water Quality Criteria

NOTES: Bold values represent detected concentrations, shaded values exceed acute or chronic criteria.

MDL is reported for non-detected constituents.

MDL = average method detection limit

B (organic) = detected in the laboratory method blank

 ${\bf J}$ (organic) = compound was detected, but below the reporting limit (value is estimated)

TABLE 26. BUTYLTIN CONCENTRATIONS (UG/L) IN SITE WATER AND STANDARD AND EFFLUENT ELUTRIATES CALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

ANALYTE	UNITS	Average RL	Site Water
DIBUTYLTIN	UG/L	0.05	0.01 U
MONOBUTYLTIN	UG/L	0.01	0.05 U
TETRABUTYLTIN	UG/L	0.012	0.0086 U
TRIBUTYLTIN	UG/L	0.0086	0.012 U
2,3,7,8-TCDD	PG/L	2.8	4.8 U

Standard Elutriate						
CCU3-BAR-	CCU3-BAR-	CCU3-DIS-				
1/2-SET	3/4-SET	SET				
2.8 U	3.3 U	2.2 U				

Effluent Elutriate						
CCU3-	CCU3-	CCU3-DIS-				
BAR-1/2-	BAR-3/4-	EFF				
3.4 U	1.7 U	3.8 U				

NOTES: RL is reported for non-detected constituents.

RL = average reporting limit

TABLE 27. TCLP IN SEDIMENT CALVERT CLIFFS NUCLEAR POWER PLANT, LUSBY, MARYLAND (JULY 2010)

MG/L MG/L MG/L MG/L MG/L	Average RL 0.0 0.0 0.0 0.0	TCLP 1.0 10.0	CCU3-BAR-1/2- SED 0.0011 U 0.0045 U	CCU3-BAR-3/4- SED 0.0011 U	CCU3-DIS- SED 0.0011 U
MG/L MG/L MG/L MG/L	0.0 0.0	1.0	0.0011 U		
MG/L MG/L MG/L	0.0			0.0011 U	0.0011.11
MG/L MG/L MG/L	0.0			0.0011 U	0.0011.11
MG/L MG/L MG/L	0.0	10.0	0.0045 U		0.0011 U
MG/L MG/L				0.0045 U	0.0045 U
MG/L MG/L					
MG/L	0.0	5.0	0.005 B	0.0022 U	0.0022 U
	0.0	100	0.18 B	0.087 B	0.22
MC/I	0.0	1.0	0.00024 U	0.00031 B	0.00048 B
MG/L	0.0	5.0	0.00084 U	0.00084 U	0.00084 U
MG/L	0.0	5.0	0.0014 U	0.0014 U	0.0014 U
MG/L	0.0	0.2	0.00008 B	0.000054 B	0.000038 U
MG/L	0.0	1.0	0.0063 B	0.0057 B	0.0095 B
MG/L	0.0	5.0	0.00058 U	0.00058 U	0.00058 U
MG/L	0.0	0.03	0.00066 U	0.00066 U	0.00066 U
MG/L	0.0	0.02	0.00039 U	0.00039 U	0.00039 U
MG/L	0.0	0.4	0.00032 U	0.00032 U	0.00032 U
MG/L	0.0	0.01	0.0004 U	0.0004 U	0.0004 U
MG/L	0.0	0.01	0.00039 U	0.00039 U	0.00039 U
MG/L	0.0	10.0	0.00037 U	0.00037 U	0.00037 U
MG/L	0.0	0.5	0.0074 U	0.0074 U	0.0074 U
MG/L	0.0	7.5	0.0037 U	0.0037 U	0.0037 U
		400			0.0076 U
					0.0087 U
					0.0027 U
MG/L	0.0	200		0.0088 U	0.0088 U
MG/L	0.0	0.1	0.00092 U	0.00092 U	0.00092 U
MG/L	0.0	0.5	0.00083 U	0.00083 U	0.00083 U
MG/L	0.0	5.0	0.0031 U	0.0031 U	0.0031 U
MG/L	0.0	2.0	0.0042 U	0.0042 U	0.0042 U
MG/L	0.0	100	0.0033 U	0.0033 U	0.0033 U
MG/L	0.0	5.0	0.0036 U	0.0036 U	0.0036 U
I					
MG/L	0.0	0.7	0.043 U	0.043 U	0.043 U
					0.038 U
					0.043 U
					0.04 U
					0.043 U
					0.021 U
					0.021 U
					0.033 U
					0.035 U
					0.052 U
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NOTES: Bold values represent detected concentrations.

RL is reported for non-detected constituents.

RL = average reporting limit

B (inorganic) = compound was detected, but below the reporting limit (value is estimated)

APPENDIX A

OYSTER SURVEY REPORT

Calvert Cliffs Oyster Population and Bottom Survey

Submitted to EA Engineering, Science and Technology



August 13, 2010



The Laboratory of Kennedy Paynter

University of Maryland

College Park, MD 20742

Introduction

A survey was conducted on August 9th, 2010 at two sites in the waters offshore of the Calvert Cliffs nuclear power plant to determine the impact of plant construction on oyster bar habitat. Figure 1 outlines the two proposed sites: Discharge Pipe (0.82 acres) and Barge Slip (4.47 acres). The survey quantified bottom type (sand, shell, mud, rock or shell hash.), amount of shell on the bottom and oyster population dynamics (size, ratio of live to dead oysters, overall population) at the two sites.

Methods

A 25m x 25m grid was overlaid on each site using GIS software and one hydraulic patent tong (1.61 m^2) grab was taken in each 25m x 25m cell. Certain cells were in water that was too shallow for the survey boat to safely sample or were too close to the existing pier, and these cells were not sampled. Figure 2 shows the patent tong grids with the sampling points at the Discharge Pipe and Barge Slip sites. The bottom type and amount of shell at each grab was recorded (shell scores range from 0 = no shell to 5= tong full of shell). The number and size of all live and dead oysters in each grab was also recorded.

<u>Results</u>

A total of 48 individual patent tong grabs were taken at the Discharge Pipe and Barge Slip sites. Of these grabs, only one contained live oysters and only three individual oysters were found in that grab.

Bottom Type

The bottom type at the survey sites was either hard bottom (sandstone, rock or shell hash) or sand. Figure 3 shows the bottom type over the survey area as either hard bottom (blue) or sand (red).

Shell Coverage

Shell coverage was low, with only 21% of the sites sampled contained shell hash. Of those cells, a moderate amount of shell was found (1-3 on a 5 point scale). Figure 4 shows the shell coverage over the survey area.

Oyster Status, Counts, Size and Density

Three live oysters and no dead oysters (boxes or gapers) were collected during the survey and they were all found in a single grab in one of the cells furthest from shore. The average shell height of the oysters collected was 99.33mm at a density of 1.66 oysters/m². Figure 5 shows the oyster density over the survey area.

Conclusions

Although very few live oysters were found in the survey area, most of the bottom surveyed in the barge slip area (4.47) could serve as oyster habitat. The discharge pipe area was all sand, which is not considered oyster habitat.

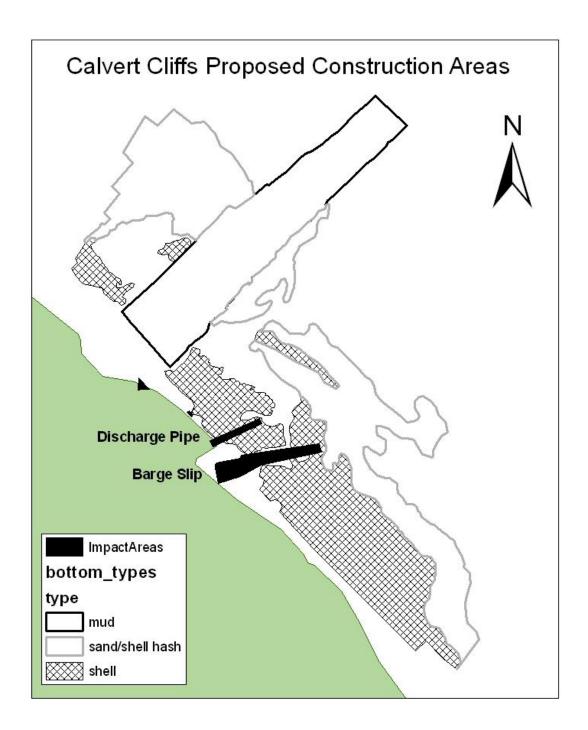


Figure 1. Areas of proposed construction at Calvert Cliffs (black polygons).

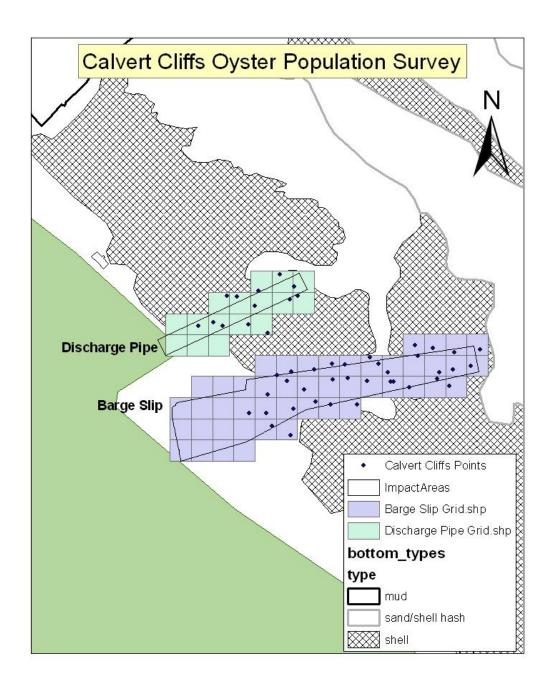


Figure 2. Patent tong survey points at Barge Slip and Discharge Pipe sites. One patent tong grab was taken at each point to quantify bottom type, shell coverage and oyster population dynamics at each proposed construction site.

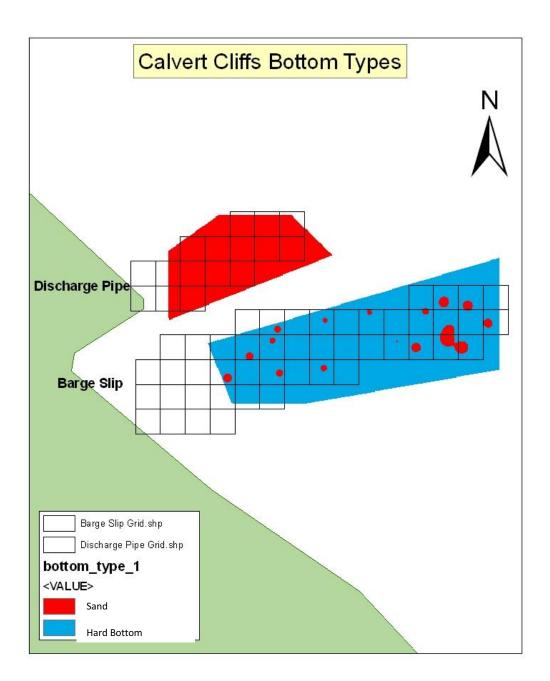


Figure 3. Bottom Type (hard bottom or sand) at Barge Slip and Discharge Pipe sites. The Discharge Pipe site was entirely sand while the barge slip site was mostly hard bottom.

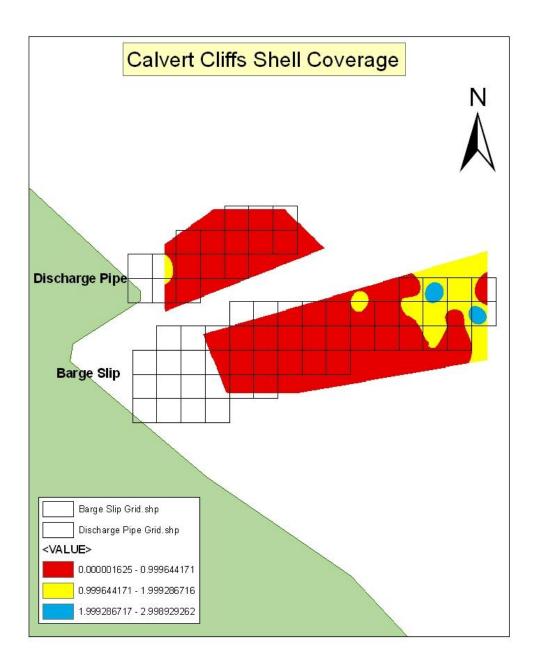


Figure 4. Shell coverage (1-5 scale with 5 being high shell coverage) at Barge Slip and Discharge Pipe sites. Almost no shell coverage was observed at the Discharge Pipe site while a moderate amount of shell coverage was observed at the northeast corner of the Barge Slip site.

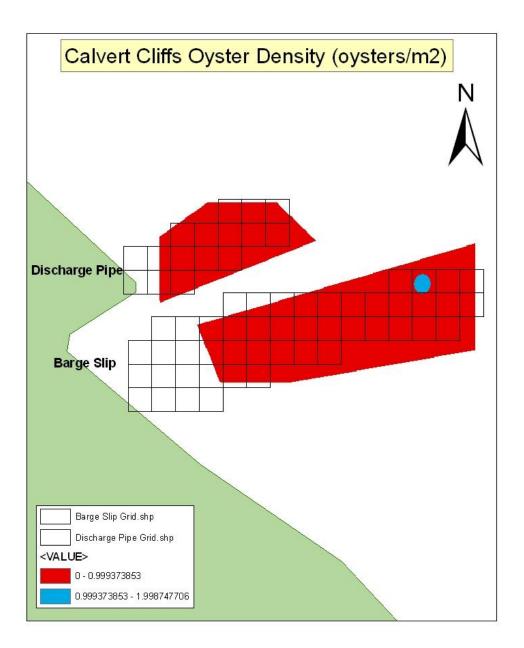


Figure 5. Oyster density (oysters/m²) at Barge Slip and Discharge Pipe sites. Oysters were only observed in one grid cell in the northeast corner of the Barge Slip site, at a density of 1.657 oysters/m²).

APPENDIX B

FIELD NOTES

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APPENDIX C

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APPENDIX D

PROJECT LIMITS

Analyte	Units	Laboratory Reporting Limit	Recommended Target Detection Limit
Metals, ITM (SW846 6020)	<u>1</u>		
Aluminum	mg/kg	2	50
Antimony	mg/kg	2	2.5
Arsenic	mg/kg	2	5.0
Barium	mg/kg	1	700^{1}
Beryllium	mg/kg	0.5	2.5
Cadmium	mg/kg	0.5	0.3
Calcium	mg/kg	10	NA ¹
Chromium	mg/kg	1.0	5.0
Cobalt	mg/kg	5.0	0.1
Copper	mg/kg	2.5	5.0
Iron	mg/kg	10	50
Lead	mg/kg	2	5.0
Magnesium	mg/kg	10	NA^1
Manganese	mg/kg	1.5	5.0
Mercury	mg/kg	0.033	0.2
Nickel	mg/kg	4.0	5.0
Potassium	mg/kg	10	NA^1
Selenium	mg/kg	2.0	1.0
Silver	mg/kg	1.0	0.2
Sodium	mg/kg	10	NA^1
Thallium	mg/kg	1.0	0.2
Tin	mg/kg	5.0	0.5
Vanadium	mg/kg	0.1	370 ¹
Zinc	mg/kg	2.0	15
Wet Chemistry			
Ammonia	mg/kg	2.4	0.1
Nitrate+Nitrite	mg/kg	20.0	
Total Kjeldahl Nitrogen	mg/kg	20.0	
Sulfide	mg/kg	4.0	0.1
Total Phosphorus	mg/kg	2.5	
TOC (Lloyd Kahn)	mg/kg	500	1000
Total Petroleum Hydrocarbons (TPH)			
Diesel Range Organics (DRO)	mg/kg	10	0.24
Gas Range Organics (GRO)	μg/kg	100	28
Chlorinated Pesticides			
Aldrin	µg/kg	1.3	10
alpha-BHC	μg/kg	1.3	
beta-BHC	μg/kg	1.3	
delta-BHC	μg/kg	1.3	
gamma-BHC (Lindane)	μg/kg	1.3	10
Chlordane (Technical)	μg/kg	33	10
Dachtal	μg/kg	TBD	2
4,4'-DDD	μg/kg	1.3	10

TABLE C-1. PROJECT LIMITS FOR SEDIMENT SAMPLESCalvert Cliffs Sediment Characterization (2010)

		Laboratory	Recommended Target Detection
Analyte	Units	Reporting Limit	Limit
4,4'-DDE	µg/kg	1.3	10
4,4'-DDT	µg/kg	1.3	10
Dieldrin	µg/kg	1.3	10
Endosulfan I	µg/kg	1.3	10
Endosulfan II	µg/kg	1.3	10
Endosulfan sulfate	µg/kg	1.3	10
Endrin	µg/kg	1.3	5
Endrin aldehyde	µg/kg	1.3	5
Heptachlor	µg/kg	1.3	10
Heptachlor epoxide	µg/kg	1.3	10
Methoxychlor	µg/kg	3.3	10
Mirex	µg/kg	1.3	
Toxaphene	µg/kg	16.5	50
Polychlorinated Biphenyl (PCB) Congeners			
2,4'-Dichlorobiphenyl (BZ # 8)	µg/kg	0.17	1
2,2',5-Trichlorobiphenyl (BZ # 18)	µg/kg	0.17	1
2,4,4'-Trichlorobiphenyl (BZ # 28)	µg/kg	0.17	1
2,2',3,5'-Tetrachlorobiphenyl (BZ # 44)	µg/kg	0.17	1
2,2',4,5'-Tetrachlorobiphenyl (BZ # 49)	µg/kg	0.17	1
2,2',5,5'-Tetrachlorobiphenyl (BZ # 52)	µg/kg	0.17	1
2,3',4,4'-Tetrachlorobiphenyl (BZ # 66)	µg/kg	0.17	1
3,3',4,4'-Tetrachlorobiphenyl (BZ # 77)	µg/kg	0.17	1
2,2',3,4,5'-Pentachlorobiphenyl (BZ # 87)	µg/kg	0.17	1
2,2',4,5,5'-Pentachlorobiphenyl (BZ # 101)	µg/kg	0.17	1
2,3,3',4,4'-Pentachlorobiphenyl (BZ # 105)	µg/kg	0.17	1
2,3',4,4',5-Pentachlorobiphenyl (BZ # 118)	µg/kg	0.17	1
3,3',4,4',5-Pentachlorobiphenyl (BZ # 126)	µg/kg	0.17	1
2,2',3,3',4,4'-Hexachlorobiphenyl (BZ # 128)	µg/kg	0.17	1
2,2',3,4,4',5'-Hexachlorobiphenyl (BZ # 138)	µg/kg	0.17	1
2,2',4,4',5,5'-Hexachlorobiphenyl (BZ # 153)	µg/kg	0.17	1
2,3,3',4,4',5-Hexachlorobiphenyl (BZ # 156)	µg/kg	0.17	1
3,3',4,4',5,5'-Hexachlorobiphenyl (BZ # 169)	µg/kg	0.17	1
2,2',3,3',4,4',5-Heptachlorobiphenyl (BZ # 170)	µg/kg	0.17	1
2,2',3,4,4',5,5'-Heptachlorobiphenyl (BZ # 180)	µg/kg	0.17	1
2,2',3,4,4',5',6-Heptachlorobiphenyl (BZ # 183)	µg/kg	0.17	1
2,2',3,4,4',6,6'-Heptachlorobiphenyl (BZ # 184)	µg/kg	0.17	1
2,2',3,4',5,5',6-Heptachlorobiphenyl (BZ # 187)	µg/kg	0.17	1
2,2',3,3',4,4',5,6-Octachlorobiphenyl (BZ # 195)	µg/kg	0.17	1
2,2',3,3',4,4',5,5',6-Nonachlorobiphenyl (BZ # 206)	µg/kg	0.17	1
2,2',3,3',4,4',5,5',6,6'-Decachlorobiphenyl (BZ # 209)	µg/kg	0.17	1
Polychlorinated Biphenyl (PCB) Aroclors		· · · · · · · · · · · · · · · · · · ·	
Aroclor 1016	ug/kg	33	
Aroclor 1221	ug/kg	33	
Aroclor 1221	ug/kg	33	
Aroclor 1242	ug/kg	33	
Aroclor 1248	ug/kg	33	
Aroclor 1254	ug/kg	33	

Analyte	Units	Laboratory Reporting Limit	Recommended Target Detection Limit
Aroclor 1260	ug/kg	33	
Polynuclear Aromatic Hydrocarbons (PAHs)			
Acenaphthene	µg/kg	6.7	20
Acenaphthylene	µg/kg	6.7	20
Anthracene	µg/kg	6.7	20
Benzo(a)anthracene	µg/kg	6.7	20
Benzo(b)fluoranthene	µg/kg	6.7	20
Benzo(k)fluoranthene	µg/kg	6.7	20
Benzo(ghi)perylene	µg/kg	6.7	20
Benzo(a)pyrene	µg/kg	6.7	20
Chrysene	µg/kg	6.7	20
Dibenzo(a,h)anthracene	µg/kg	6.7	20
Fluoranthene	µg/kg	6.7	20
Fluorene	µg/kg	6.7	20
Indeno(1,2,3-cd)pyrene	µg/kg	6.7	20
2-Methylnaphthalene	µg/kg	6.7	20
1-Methylnaphthalene	µg/kg	6.7	20
Naphthalene	µg/kg	6.7	20
Phenanthrene	µg/kg	6.7	20
Pyrene	µg/kg	6.7	20
Dioxins			
2,3,7,8-TCDD	µg/kg	1	1
Butyltins			
Dibutyiltin	µg/kg	10.0	10
Monobutyltin	µg/kg	10.0	10
Tetrabutyltin	µg/kg	10.0	10
Tributyltin	µg/kg	10.0	10

¹ Analytes added for Shirley Plantation screening. Target detection limit is based on the Shirley Plantation Screening Criteria. NA values are values for which Shirley Plantation has not provided a screening value.

Calvert Cliffs Se	ediment Ch	aracterization (20	
Analyte	Units	Laboratory Reporting Limit	Target Detection Limit
Volatiles, TCLP (SW846 1311/8260E	<u>3)</u>		
Benzene	μg/L	0.050	0.50
2-Butanone	μg/L	0.050	200
Carbon tetrachloride	μg/L	0.050	0.50
Chlorobenzene	μg/L	0.050	100
Chloroform	μg/L	0.050	6.0
1,2-Dichloroethane	μg/L	0.050	0.50
1,1-Dichloroethene	μg/L	0.050	0.70
Tetrachloroethene	μg/L	0.050	0.50
Trichloroethene	μg/L	0.050	0.70
Vinyl chloride	μg/L	0.050	0.20
Semivolatiles, TCLP (SW846 1311/8	270C)		
Cresols (total)	μg/L	0.050	200
1,4-Dichlorobenzene	μg/L	0.010	7.5
2,4-Dinitrotoluene	μg/L	0.050	0.13
Hexachlorobenzene	μg/L	0.010	0.13
Hexachlorobutadiene	μg/L	0.01	0.50
Hexachloroethane	μg/L	0.050	3.0
3-Methylphenol & 4-Methylphenol	μg/L		
Nitrobenzene	μg/L	0.01	2.0
Pentachlorophenol	μg/L	0.05	100
Pyridine	μg/L	0.05	5.0
2,4,5-Trichlorophenol	μg/L	0.050	400
2,4,6-Trichlorophenol	μg/L	0.050	2.0
Pesticides, TCLP (SW846 8081A)			
Gamma- BHC (Lindane)	μg/L	0.0005	0.40
Chlordane (technical)	μg/L	0.005	0.030
Endrin	μg/L	0.0005	0.20
Heptachlor	μg/L	0.0005	0.0080
Heptachlor epoxide	μg/L	0.0005	0.0080
Methoxychlor	μg/L	0.001	10
Toxaphene	μg/L	0.02	0.50
Herbicides, TCLP (SW846 1311/815	1A)		
2,4-D	μg/L	0.04	10
2,4,5-TP (Silvex)	μg/L	0.01	1.0
Metals, TCLP (SW846 6010B/7470A	()		
Arsenic	μg/L	0.50	5.0
Barium	μg/L	10	100
Cadmium	μg/L	0.10	1.0
Chromium	μg/L	0.50	5.0
Lead	μg/L	0.5	5.0
Mercury	μg/L	0.0002	0.2
Selenium	μg/L	0.25	1.0
Silver	μg/L	0.25	5.0

TABLE C-2. PROJECT LIMITS FOR TOXICITY CHARACTERISTIC LEACHING PROCEDURE (TCLP) SAMPLES Calvert Cliffs Sediment Characterization (2010)

Analyte	Units	Laboratory Reporting Limit	Recommended Target Detection Limit
Metals, ITM (SW846 6020)	<u> </u>		
Aluminum	μg/L	100	40
Antimony	μg/L	4.0	3
Arsenic	μg/L	2.0	1
Barium	μg/L	10	NA^1
Beryllium	μg/L	2.0	0.2
Cadmium	μg/L	2.0	1
Calcium	μg/L	100	NA^1
Chromium	μg/L	10.0	1
Cobalt	μg/L	2.0	4
Copper	μg/L	10.0	1
Iron	μg/L	100	10
Lead	μg/L	1.0	1
Magnesium	μg/L	100	NA^1
Manganese	μg/L	2.0	1
Mercury	μg/L	0.2	0.2
Nickel	μg/L	10.0	1
Potassium	μg/L	100	NA^1
Selenium	μg/L	10.0	1
Silver	μg/L	10.0	1
Sodium	μg/L	100	NA ¹
Thallium	μg/L	1.0	1
Tin	μg/L	10.0	5
Vanadium	μg/L	1	NA ¹
Zinc	μg/L	10.0	1
Wet Chemistry		• •	
Ammonia	mg/L	0.2	0.03
Nitrate+Nitrite	mg/L	0.1	
Total Kjeldahl Nitrogen	mg/L	0.2	
Sulfide	mg/L	0.2	
Total Phosphorus	mg/L	0.050	
TOC (Lloyd Kahn)	mg/L	1	0.1
Total Petroleum Hydrocarbons (THP)	·		
Diesel Range Organics (DRO)	μg/L	100	47
Gas Range Organics (GRO)	μg/L	100	28
Chlorinated Pesticides		· · · · ·	
Aldrin	µg/L	0.02	0.04
alpha-BHC	μg/L	0.02	
beta-BHC	μg/L	0.02	
delta-BHC	μg/L	0.02	
Gamma-BHC (Lindane)	μg/L	0.02	0.1
Chlordane (Technical)	μg/L	0.5	0.17

TABLE C-3. PROJECT LIMITS FOR SITE WATER AND ELUTRIATE SAMPLES Calvert Cliffs Sediment Characterization (2010)

A se a la da	The Ar	Laboratory	Recommended Target Detection
Analyte		Reporting Limit	Limit
Dachtal	μg/L	TBD 0.02	0.03
4,4'-DDD	μg/L		0.01
4,4'-DDE	μg/L	0.02	0.01
4,4'-DDT	μg/L	0.02	0.01
Dieldrin	μg/L	0.02	0.02
Endosulfan I	μg/L	0.02	0.1
Endosulfan II	μg/L	0.02	0.1
Endosulfan sulfate	μg/L	0.02	0.1
Endrin	μg/L	0.02	0.1
Endrin aldehyde	μg/L	0.02	0.1
Heptachlor	μg/L	0.02	0.1
Heptachlor epoxide	μg/L	0.02	0.1
Methoxychlor	μg/L	0.05	0.5
Mirex	μg/L	0.02	
Toxaphene	μg/L	0.25	0.5
Polychlorinated Biphenyl (PCB) Congeners			
2,4'-Dichlorobiphenyl (BZ # 8)	ng/L	10	8.272
2,2',5-Trichlorobiphenyl (BZ # 18)	ng/L	1	0.48
2,4,4'-Trichlorobiphenyl (BZ # 28)	ng/L	1	0.432
2,2',3,5'-Tetrachlorobiphenyl (BZ # 44)	ng/L	1	0.436
2,2',4,5'-Tetrachlorobiphenyl (BZ # 49)	ng/L	1	0.449
2,2',5,5'-Tetrachlorobiphenyl (BZ # 52)	ng/L	1	0.431
2,3',4,4'-Tetrachlorobiphenyl (BZ # 66)	ng/L	1	0.505
3,3',4,4'-Tetrachlorobiphenyl (BZ # 77)	ng/L	1	0.441
2,2',3,4,5'-Pentachlorobiphenyl (BZ # 87)	ng/L	1	0.407
2,2',4,5,5'-Pentachlorobiphenyl (BZ # 101)	ng/L	1	0.413
2,3,3',4,4'-Pentachlorobiphenyl (BZ # 105)	ng/L	1	0.383
2,3',4,4',5-Pentachlorobiphenyl (BZ # 118)	ng/L	1	0.532
3,3',4,4',5-Pentachlorobiphenyl (BZ # 126)	ng/L	1	0.393
2,2',3,3',4,4'-Hexachlorobiphenyl (BZ # 128)	ng/L	1	0.356
2,2',3,4,4',5'-Hexachlorobiphenyl (BZ # 138)	ng/L	1	0.338
2,2',4,4',5,5'-Hexachlorobiphenyl (BZ # 153)	ng/L	1	0.392
2,3,3',4,4',5-Hexachlorobiphenyl (BZ # 156)	ng/L	1	0.374
3,3',4,4',5,5'-Hexachlorobiphenyl (BZ # 169)	ng/L	1	0.429
2,2',3,3',4,4',5-Heptachlorobiphenyl (BZ # 170)	ng/L	1	0.368
2,2',3,4,4',5,5'-Heptachlorobiphenyl (BZ # 170)	ng/L	1	0.364
2,2',3,4,4',5',6-Heptachlorobiphenyl (BZ # 180)	ng/L	1	0.372
2,2',3,4,4',6,6'-Heptachlorobiphenyl (BZ # 183)	ng/L	1	0.423
2,2',3,4',5,5',6-Heptachlorobiphenyl (BZ # 184)	ng/L	1	0.394
2,2,3,3,4,4,5,6-Octachlorobiphenyl (BZ # 187)	ng/L	1	0.393
	ng/L ng/L	1	0.383
2,2',3,3',4,4',5,5',6-Nonachlorobiphenyl (BZ # 206) 2,2',3,3',4,4',5,5',6,6'-Decachlorobiphenyl (BZ # 209)	ng/L ng/L	1	0.438

Polychlorinated Biphenyl (PCB) Aroclors			
Aroclor 1016	μg/L	1	
Aroclor 1221	μg/L	1	
Aroclor 1232	μg/L	1	
Aroclor 1242	μg/L	1	
Aroclor 1248	μg/L	1	
Aroclor 1254	μg/L	1	
Aroclor 1260	μg/L	1	
Polynuclear Aromatic Hydrocarbons (PAHs)			
Acenaphthene	μg/L	0.2	10
Acenaphthylene	μg/L	0.2	10
Anthracene	μg/L	0.2	10
Benzo(a)anthracene	μg/L	0.2	10
Benzo(b)fluoranthene	μg/L	0.2	10
Benzo(k)fluoranthene	μg/L	0.2	10
Benzo(ghi)perylene	μg/L	0.2	10
Benzo(a)pyrene	μg/L	0.2	10
Chrysene	μg/L	0.2	10
Dibenzo(a,h)anthracene	μg/L	0.2	10
Fluoranthene	μg/L	0.2	10
Fluorene	μg/L	0.2	10
Indeno(1,2,3-cd)pyrene	μg/L	0.2	10
2-Methylnaphthalene	μg/L	0.2	10
1-Methylnaphthalene	μg/L	0.2	10
Naphthalene	μg/L	0.2	10
Phenanthrene	μg/L	0.2	10
Pyrene	μg/L	0.2	10
Dioxins			
2,3,7,8-TCDD	μg/L	10	10

¹ Analytes added to metals listed based on Shirley Plantation Screening Table, which does not use aqueous results.