Attachment E

Engineering Contingencies

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<center>**Attachment E**</center>

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1.0   Introduction

This attachment describes engineering contingencies that may be applied in the event that action levels or the Resuspension Standard threshold are exceeded. The levels of the performance standard were developed using statistical analysis of historical data, surface water quality modeling and applicable federal standards. The resuspension criteria will be used to implement various engineering contingencies to minimize the release of PCBs during the remediation and to achieve the remediation goals as set forth in the Record of Decision (ROD) (USEPA, 2002). In the event that the resuspension criteria are exceeded, engineering contingencies will be implemented as necessary to minimize the potential impact of dredging on ambient water quality. A series of contingencies, ranging from increased monitoring frequency to cessation of dredging operation, have been proposed. These engineering contingencies will be implemented based on near-field and far-field water quality monitoring results.

The performance standard requires additional monitoring under certain conditions. The frequency and parameters for this additional monitoring are defined as a part of the performance standard. For other contingencies (*i.e.*, contingencies not specifically addressed in the performance standard), the specific technology cannot be selected, but must be a judgment that is specific to the problem encountered. Contingencies must be developed during the design stage for use in the event that water column concentrations exceed the performance standard. The performance standard does specify that if certain levels are exceeded, the cause of the exceedance will be examined and necessary changes must be made to the existing operations.

This attachment provides a brief overview of the performance standard (including a discussion of the monitoring locations needed to assess compliance with the standard), a summary of engineering contingencies used during similar projects, and a discussion of the engineering contingencies that may be applicable to the remediation.

Engineering contingencies for the Public Water Intakes and agricultural water intakes will be addressed in the Community Health and Safety Plan.

1.1   Performance Standard Monitoring Locations

Two types of monitoring locations are discussed throughout this attachment. Definitions are provided below:

Far-Field (Upper River and Lower River)

Far-field stations are fixed locations, typically located at dams and bridges. The primary contaminants to be monitored at these stations are PCBs and suspended solids. The results from monitoring at the far-field stations are the primary measure of PCB loss due to dredging, based

on the assumption that only PCBs escaping each river section have the potential to cause significant downstream impacts.

Near-Field

Near-field monitoring locations are located within a short range of the remedial operations, typically within a mile or so downstream. Depending upon the proximity of the various ongoing remedial operations to one another and the use of barriers, each remedial operation may have near-field monitoring locations associated with it. These near-field stations will be monitored continuously to determine the local impacts. The primary measurements in the near-field will be suspended solids concentrations and turbidity.

1.2   Resuspension Criteria

The resuspension criteria consist of three action levels and one standard providing limits on PCB and suspended solids concentration. Each of the resuspension criteria has associated monitoring requirements and engineering contingencies. The monitoring plan is summarized in Tables 1-2, 1-3 and 1-4 of the main document, showing the parameters required at each station and the frequency of sampling. Table 1-1 of the main document lists the concentration or load limits for each action level. Monitoring and resuspension criteria are fully described in the main body of the text and in Attachment F. An engineering evaluation of conditions in the river leading to elevated concentrations is recommended for Evaluation Level, but is mandatory for the Concern Level, Control Level and Resuspension Standard threshold. Similarly, implementation of engineering contingencies to reduce contaminant levels in the river is recommended at Evaluation Level, but is mandatory for other the three other levels.

2.0   Monitoring and Contaminant Control Technologies Used At Other Sites

The monitoring and contaminant control technologies employed at three other PCB remediation sites are described below. The three sites are:

- St. Lawrence River Remediation Project at the Alcoa, Inc. Massena East Smelter Plant, New York, (Bechtal Environmental, 2000; 2002)

- New Bedford Harbor (Pre-Design Field Test), New Bedford, Massachusetts, (USACE, August, 2001); and

- Grand Calumet River, Gary, Indiana, (Earth Tech, Inc., 2002).

The technologies implemented at these three sites and reviewed in this attachment are containment (St. Lawrence River), dredging system design [hydraulic bucket design] (New Bedford Harbor), and monitoring (Grand Calumet River).

2.1 St. Lawrence River Remediation Project at the Alcoa, Inc. Massena East Smelter Plant, New York (Reynolds Metals)

In order to control export of PCB-contaminated sediment at the St. Lawrence River Alcoa site, a containment system was installed as part of the remedial design. The containment system at this site included:

- a sheet pile wall that enclosed the entire remediation area;

- silt curtains that provided secondary containment for the more highly contaminated Area C and also isolated uncontaminated portions of Area B from dredging areas; and

- air gates (air curtain technology) that created air-bubble curtain that acted as a circulation barrier while allowing for barge and tugboat access to areas enclosed by the silt curtain and pile wall.

Each of these components is discussed below.

Sheet Pile Wall

The wall consisted of interlocking steel sheeting embedded several feet or more into sediments and supported by H-beams ("king piles") driven to greater depths. The sheeting and king piles were tied together through a welded and bolted framework of steel braces and walers. The 3,800-feet finished wall consisted of about 200 king piles and 2,200 sheets. The maximum depth of water along the wall was about 32 feet.

The original design of the sheet pile wall specified that every fifth sheet would be driven to the water surface to balance any differences in hydrostatic pressure between the inside wall and the outside. However, this was later changed and all sheets were raised to a height of about 2 ft above the river surface, minimizing the connection of turbid water inside the sheet pile wall with the river water outside the enclosure.

After the installation, a video survey was conducted to verify that there were no openings along the bottom of the wall or open seams in the sheeting. This survey identified a few small holes that were patched using sand bags. In addition, some of the sheeting was trimmed to get all the sheets down to the 2 ft above water level after installation to reduce the surface area exposed to wind forces. Environmental monitoring data showed that the sheet pile wall functioned as designed and effectively contained the turbidity and suspended sediments generated during the dredging activities within the remediation area.

Silt Curtains

Silt curtains, consisting of 22-oz. PVC sheeting weighted on the bottom and suspended by polystyrene floatation buoys, were installed around Area C and a portion of Area B. The silt curtains were tied to H-beam anchor posts driven at a spacing of 100 feet, and anchored on the shoreline of a driven post or tree. The ballast for the curtains was 3/8-in. galvanized anchor

chain within a sealed pocket in the sheeting that could adapt to the bottom contours, thereby providing a complete vertical barrier. The curtain was suspended by cables attached to tensioners and anchor plates with reefing lines connected to the lower ballast chain to adjust the vertical height. A total of 1,222 feet and 996 feet silt curtains were used in Area B and Area C, respectively. The silt curtains effectively isolated the more contaminated Area C and prevented contamination of the clean portion of Area B.

The original design called for the installation of the silt curtain H-beam piles after the sheet pile wall was completed. However, due to additional time required installing the sheet pile wall, this plan was changed for the clean part of Area B, and the silt curtain H-beam piles were driven while the sheet pile wall was being installed. A similar change for the contaminated part of Area B was not approved by USEPA.

Another change to the design of the silt curtain involved the addition of dual H-beams rather than a single H-beam to anchor the curtain. The original design specified that one H-beam would be placed at intervals along the inside of the curtain and timbers would be attached to the top of the beam to prevent barge traffic from hitting the curtain from outside. The silt curtain manufacturer recommended placing dual H-beams at a spacing of 90 feet and then anchoring the curtain between the beams.

Air Gates

Air gates (air curtain technology) were used to create vertical circulation barriers that allow boats to pass but restrict the movement of water between various parts of the remediation area. The air curtains consisted of 2-in. outside diameter (OD) steel pipe fitted with diffuser orifices on a helical, 9-inch spacing. The pipes had leg supports that raised them about a foot off the bottom. Geomembrane was laid beneath the pipes to minimize the disturbance of nearby sediment. Divers were used to place the liner, pipe and anchors, connect the supply lines and verify proper operation once the equipment was in place. A compressor station supplied air to the gates at a flow rate of about 1,000 cubic feet per minute (cfm) with flow pressures of 90 to 100 psig. The gates allowed for barge transit and limited the migration of turbid water across the barrier. A major objective of the gates was to contain the turbidity generated during the removal of Area C sediment. The gates accomplished this objective and otherwise functioned as designated for the duration of the project.

2.2 New Bedford Harbor (Pre-Design Field Test), New Bedford, Massachusetts

A pre-design field test was conducted at the New Bedford Harbor site to assess the effectiveness of hydraulic dredging as an engineering contingency to minimize the release of PCB contaminated material to the water column and to limit the transport of sediment away from the dredging area. The water quality monitoring data obtained during dredging activities indicated that the actual dredging process using hydraulic excavator appeared to have a limited impact on water column. The factors that minimized.the release of material to the water column included the design of the bucket (tight closing with limited leakage), the configuration of the dredge (with a "moon-pool" work area enclosed behind a 36-inch silt curtain) and the controlled manner in which the operation was executed.

Factors that limited the transport of contaminated material away from the dredging area included the shallowness of the area (maximum depth of the dredged area was less than 10 feet (3 m) at high tide and the limited currents (maximum currents generally less than 0.5 feet/sec.).

Activities performed in support of the dredging (operation of support vessels such as tug boats) appeared to have a much greater impact on water quality than the dredging operations due to shallowness of the water (about 4 to 5 feet).

Normal fluctuations that occur in Upper Harbor due to changing environmental conditions appeared to be similar or greater in scale than the overall impacts related to the actual dredging process.

2.3   Grand Calumet River, Gary, Indiana

Dredging activities are scheduled for completion in December 2003. The following was extracted from the Water Quality Certification Work Plan dated July 2002.

Three water quality monitoring locations (Sites A, B, and C) are defined as the primary monitoring sites. A fourth monitoring location (Site D) is defined as the verification site.

- Site A is located to monitor water quality upstream of dredging (located mid-channel of the Grand Calumet River at Transect 4 and will be re-located to Transect 2 as dredging progresses),

- Site B is located mid-channel, approximately 200 yards downstream of the open water dredge in Transect 12 to 36, and will be re-located as dredging progresses through cell D (or from Transect 12 to 36),

- The third station, Site C, is the downstream sample site and is located mid-channel downstream of Transect 36 (downstream of the limit of dredging), and

- A fourth sample location, Site D, also known as the verification sample site, will be situated 200 yards upstream of the open water dredge in transects 12 to 36 and will be used to verify water quality exceedances and used to determine if the exceedance is a result of the dredging operation or a different point source. This station was proposed instead of performing background sampling prior to initiating dredging. All water samples will be equal volume composites created from a total of three samples per location. These three samples per location will be taken from the water surface, at 50 percent of the water depth and at 80 percent of the water depth.

Three levels of monitoring will be utilized, which includes Level 3 Monitoring (*i.e.*, collection of composite water samples once per month from automatic samplers at Sites A and C and manually at Sites B and D for analysis of PCBs and other specified parameters). If results indicate no exceedances at Sites A, B and C or if monitoring indicates exceedances at all three sites (A, B, and C), then it will be concluded that dredging is not the source and normal sampling

will be conducted (once per month). If, however, results indicate exceedances at Sites B and C but not site A, then the water sample collected at Site D will be analyzed. If the sample from Site D indicates the parameters exceeded at Sites B and C are also exceeded at Site D, it will be assumed that the downstream exceedances at these sites are not a result of dredging and the normal frequency sample will be conducted. However, if no exceedances are found at Site D, it will be concluded that dredging is the source and enhanced monitoring consisting of additional sample collection at Sites A, B and C will be implemented at a rate of three times per week. When results indicate that the parameters of concern are less than the criterion for two months of consecutive samples, enhanced monitoring will be discontinued and the normal monitoring frequency will be resumed.

In addition to the increased sampling frequencies as a result of exceedances determined to be due to dredging, the Indiana Department of Environmental Management and the US Army Corps of Engineers will also implement a response action. If it is thought that an immediate threat to human health or aquatic life exist, the required response action will be issued within 72 hours and this action will be implemented as quickly as possible with a maximum time limit of implementation of one week. If this schedule is not met, enhanced monitoring will be automatically implemented as described above, based on the parameters exceeded and the level of monitoring utilized when the exceedances occurred.

Possible response actions may consist of the following engineering contingencies:

- Decrease dredging operation,

- Install additional turbidity barriers or control mechanisms,

- Temporary cessation of dredging activities, and

- Conduct additional monitoring.

3.0 Engineering Contingencies for the Remediation

The required engineering contingencies for the Resuspension Performance Standard are described below. These include increased monitoring frequency, engineering studies, containment technologies, operational modifications, equipment modifications and scheduling changes. With the exception of monitoring frequency, specific implementations of the engineering contingencies must be planned during design.

The applicability of many of the containment technologies was evaluated in the Appendix E.5 of the FS (USEPA, 2000). The advantage and limitation of each turbidity barrier were discussed. This information will be useful when choosing the appropriate containment system for a specific area to address the engineering contingency during the remediation.

3.1 Monitoring Contingencies

Monitoring frequency of the far-field stations will be increased at higher levels of exceedance to gain more information from which to evaluate conditions. The degree of increased frequency is detailed in Table 1-2, 1-3 and 1-4 of the main document for non-routine monitoring. The sampling method also changes for some stations -- from grab samples to composites of hourly samples -- to better capture the average water column concentration at the nearest representative far-field stations and to limit the number of analytical samples required.

3.2 Engineering Evaluations

In instances where water quality measurements exceed a resuspension criteria based on PCB concentration or load, an evaluation of the remedial operations should be conducted to determine the possible source and mechanism causing the exceedance, including:

- Examine the barrier, if it is in use, for leaks and stability,

- Examine the sediment transport pipeline if a hydraulic dredge is used,

- Examine the turbidity associated with sediment transport barges and other support vehicles, and

- Sample PCB concentrations in the near-field.

These engineering studies will be mandatory for exceedance of the Concern Level, Control Level and Resuspension Standard threshold.

3.3 Barriers

Several types of barrier systems are described below:

- Fixed Structural Barriers,

- Non-Structural (Portable) Barriers,

- Other Portable Barrier Systems, and

- Control Zone Technology.

Fixed Structural Barriers

Fixed structural barriers such as sheet piling are particularly suitable for areas where potential for high levels of resuspension are expected. Sheet piling consists of a series of interlocking steel sections. The piles are all driven in panels to approximately the same depth. It is not anticipated that turbidity barriers comprised of sheet piling will have applicability to areas where relatively shallow rock is present.

While fixed structural barriers provide considerable structural capacity, these systems are relatively expensive and usually require significant planning, equipment and manpower resources to install.

Non-Structural Barriers (Portable Barriers)

Non-structural barriers, such as silt curtains and silt screens (sediment curtains), can be considered for use to contain the sediment transport during dredging. Silt curtains are constructed of impervious materials that block or deflect the passage of water and sediments. Silt screens are similar to silt curtains; however, these barriers allow water to flow through while impeding the passage of a fraction of the suspended load. Typically, a silt curtain and silt screen are suspended by a flotation unit at the water surface and held in a vertical position by a ballast chain within the lower hem of the skirt. Anchors attached to the barrier also serve to hold it in place.

The advantage of using non-structural barriers is that they can easily be deployed and re-located to new work areas after dredging at a specific location has been completed. Silt curtains are not considered appropriate in situations where the river current is greater than approximately 1.5 feet per second and where the depth of the river exceeds 21 feet. However, it should be noted that if the silt curtain is set up in a configuration that is closely parallel to river flow, the curtain could function effectively in currents approaching 3 feet per second.

Other Portable Barrier Systems

Other commercial products such as the Portadam^{TM} and Aqua-Barrier^{TM} systems are also available for construction site containment, diversion of water flow, erosion control and flood control. These systems are low-cost alternatives to building earthen dams or using sheet piles, and are relatively easy to set up. These systems are generally applicable to water depths of less than 10 feet.

The Portadam^{TM} system utilizes a freestanding steel support structure in conjunction with an impervious fabric membrane. The support members transfer fluid loading to an approximately vertical downward load, allowing for installation on a solid impenetrable foundation. This structure free-stands on the existing bed, which eliminates the need for pile-driving equipment, cross bracing or anchorage. The membrane is placed on the outer section of the support structure, and is rolled out all the way down to the level of the bed. Hydraulic loading on the membrane assists in the sealing and stability of the entire structure. Once installed, the work area enclosed by the structure can be de-watered.

The Aqua-Barrier[TM] and GeoCHEM Water Structures[TM] systems utilize water-filled vinyl polyester-reinforced tubes to provide mass for stability and they can be coupled together to form a barrier of any length. Punctures in the material may be easily patched with repair kits. They are lightweight, easy to transport and re-usable. While these systems are not as sturdy as the Portadam[TM] system, they can be used in cold weather conditions and are reasonably resistant to sunlight exposure.

Air gates are used to facilitate the passage of dredging related traffic to and from an enclosed (i.e., sheet piled or silt curtained) area. The technology employs a continuous release of bubbles to reduce the flow of water to and from an enclosure. The air is supplied from a blower or compressed air source. The effort and cost associated with the deployment and operation of air gates are low and the performance of air gates appear to be superior compared to silt curtain gates.

Control Zone Technology

Control zone is a secure dredging area that is maintained and sealed off to prevent the release of contaminants generated inside the zone. Application of control zone technology (CZT) allows excavation of contaminated sediments without the release of particulate and soluble contaminants into the surrounding water environment. It also establishes an area that can be easily monitored to confirm that remediation goals are met. This type of technology is more stringent than other barrier technology, since it requires additional water treatment. CZT has only been tested on a pilot scale and the cost is likely to be prohibitive. This type of technology could be considered for limited use in the most highly contaminated areas.

3.4   Operation and Equipment Modifications

Depending on the level of resuspension observed, operational control and equipment modification should be considered, which include:

- Limiting boat speeds to reduce prop wash,

- Restricting the size of boats that can be used in certain areas,

- Loading barges to less than their capacity, where it is necessary to reduce draft,

- Selecting an alternate dredge with a lower resuspension rate,

- Selecting alternate equipment or method for placing backfill or capping material, and

- Use of smaller, shallow draft boats to transport crewmembers and for inspection of dredges.

3.5 Scheduling Changes

In May and June, the baseline water column concentrations are high relative compared to the remainder of the dredging season. As documented in the baseline water column level study (Attachment A), the 95 percent upper confidence limits (95% UCL) on the mean of PCB concentration at the TI Dam and Schuylerville ranged from 110 ng/L to 200 ng/L in May and June. Remedial activities in high-concentration areas during high flow conditions may result in increased water-column PCB concentrations above resuspension criteria and therefore necessitate implementation of engineering contingencies such as containment systems capable of containing enough of the resuspended material to maintain acceptable water column concentrations. Areas with higher sediment concentration may need to be scheduled for remediation in later months of each year (i.e., low flow conditions, when the baseline level of PCB concentration is relatively low) if the engineering contingencies chosen are not effective. Baseline water column concentrations should also be considered when scheduling remediation in areas nearest the WTPs in order to maintain a margin of safety for the public water supply.

4.0 Implementation Strategies

Flowcharts depicting the implementation of the Resuspension Performance Standard are provided in Figures 3-1, 3-2 and 3-3 of the main document for the near-field suspended solids, far-field total PCBs and far-field suspended solids. These flowcharts present the interaction between the three aspects of the Resuspension Performance Standard: resuspension criteria, monitoring requirements and engineering contingency requirements.

5.0 References

Bechtel Environmental, Inc./Metcalf & Eddy, Inc. 2000. Final Dredging Program Design Report for the River Remediation Project at the Reynolds Metals Company, St. Lawrence Reduction Plant, Massena, New York, Revision 3. Prepared for Reynolds Metals Company. May 2000.

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Attachment F

Measurement Technologies

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LIST OF FIGURES

APPENDIX

Attachment F

Measurement Technologies

1.0   Introduction

This section provides detailed descriptions of specific measurement techniques for the general continuous monitors prescribed in the performance standard. These include:

- o   In-situ Turbidity Measurement,

- o   In-situ Total Suspended Sediment Measurement,

- o   Semipermeable Membrane Devices (SPMDs),

- o   Trace Organic Platform Sampler (TOPS), and

- o   ISCO Portable Water And Wastewater Sampler.

These instruments are presented as examples of technology that may be used during construction to satisfy the requirements of the standard, but the selection of appropriate technology will be a part of the design process.

Several other issues related to the monitoring are presented in this attachment. Correlations between turbidity and suspended solids measurements are discussed. Development of a correlation between these parameters will be required in order to have a real time indication of dredge-related impacts on the water column. Attachment F-1 presents the results of a literature search on this topic. Attachment F-2 provides a synopsis of PCB analytical methods and associated detection limits. The detection limits for PCB congener analysis will be low in order to have detections at each station and to allow for identification in congener patterns.

2.0   Measurement Techniques

All types of dredging (navigational and environmental remedial action) create sediment plumes in the water column. Of particular interest for the Hudson River remedial action are plumes associated with (1) the mechanical and/or hydraulic dredging (sediment removal) operation, (2) material handling of dredged materials, (3) boat and barge movements, and (4) open-water placement of backfill materials. The regulatory agencies and the public are concerned about potential adverse effects caused by these plumes on humans and biological resources either through impact to water quality or increased siltation. To gain a better understanding of the temporal and spatial dynamics of sediment plumes, and, in order to implement the performance standard for resuspension, it is necessary to monitor the plumes to determine their composition, extent, and duration. Numerous techniques have been used to monitor sediment plumes, ranging from collection of water samples using simple water samplers to highly complex systems involving state-of-the-art instrumentation. Given the variety of techniques available to monitor dredging-related plumes, it is necessary to understand the advantages and limitations of the

various techniques in order to determine the techniques that provide the most cost-effective approach for particular monitoring requirements.

The resuspension performance standard (as defined in Section 1) includes specifications for both PCBs and suspended sediment. The PCB standard requires measurement of total PCB concentration, i.e. both dissolved phase and suspended phase measurements of all PCB congeners (monochlorbiphenyls through decachlorobiphenyl). Suspended solids (sediment) standards have been defined in order to serve as a surrogate for the amount of PCBs in the water column in order to provide a real-time indication of PCB concentration. Turbidity may be used as a surrogate for PCBs. Although turbidity has been historically used to estimate suspended sediment in monitoring dredging projects using empirical correlations, it is well recognized that these calibrations are site-specific and subject to significant error.

The objectives of measuring the various water quality parameters discussed above (PCBs, suspended solids, and turbidity) are twofold: first, to determine the water quality associated with the plume; and second, to track the plume both in space and time. Knowledge of the spatial extent of a given plume is necessary to determine areas of potential plume impact. Similarly, knowledge of the time history of a plume provides information on how long a plume is present in a particular area and the time required for the plume to dissipate. It is clear that both near-field and far-field monitoring are necessary.

It may be important to measure various physical parameters not directly associated with water quality such as currents, waves, and water elevations. Currents carry plumes from the area in which they were generated into adjacent waters. Therefore, data on the current structure can be used to estimate the movement and spatial extent of the plume. Waves increase turbulence in the water column that can put additional sediment into suspension and prevent material in suspension from settling out.

Measurement techniques for monitoring plumes involve either (1) collection of water samples from the water column for analysis either in the field or the laboratory (ex-situ methods), or (2) placement of instruments in the water column to directly measure water quality parameters or other physical parameters (in-situ methods). Offsite laboratory analysis is time-consuming, expensive, and cannot provide data in the short term (i.e., within a few hours or less of sample collection). At present, there are no in-situ methods available for directly measuring PCB congeners in the water column, therefore, sample collection and laboratory analysis are required.

Concentrations of PCBs in the water column are often present at parts-per-billion (μ/L) or parts-per-trillion (ng/L) levels. Conventional sampling, extraction, and analysis methods like liquid-liquid extraction or solid-phase extraction can require sampling and processing very large volumes of water (e.g., 50 liters) for analysis of adequate sensitivity to detect low concentrations. (See Attachment F-2 for a synopsis of PCB analytical methods and associated detection limits.) These limitations in methods for the direct measurement of contaminant water concentrations have often prompted the use of biomonitoring organisms for assessing the exposure of these organisms to trace/ultra-trace levels of hydrophobic chemicals like PCBs. Because certain organisms often bioconcentrate these seemingly innocuous levels of PCBs to relatively higher levels (parts per million) in their lipids, determination of the bioavailable portion of

environmental pollutants like PCBs is critical to assessing the potential for detrimental biological impacts. This organism-based approach also has inherent problems, including biotransformation and depuration of contaminants, and inapplicability in many exposure situations due to the effects of stress on the biomonitoring organisms that often lead to a lack of proportionality between the biomonitoring organism tissue concentrations and ambient exposure concentrations (Petty et al., 2000). Therefore, innovative approaches for sampling and analyzing trace/ultra-trace levels of water-borne PCBs are needed.

The major mechanisms accounting for relatively high concentrations of PCBs in organisms are the passive processes biomembrane diffusion and partitioning between an organism's lipids and its environment. Employing a mimetic chemistry approach (i.e., use of processes in simple or uniform media to mimic complex biological systems), scientists at the USGS's Columbia Environmental Research Center (CERC) have developed a passive, integrative sampler that simulates hydrophobic chemical bioconcentration. The uncertainty of estimating ambient exposure concentrations from tissue concentrations in biomonitoring organisms is thereby avoided. This sampler, the semipermeable membrane device (SPMD), measures the concentration of dissolved phase PCBs in the water column. A second type of integrating sampler has been developed by New York State Department of Environmental Conservation (NYSDEC). The Trace Organic Platform Sampler (TOPS) concentrates hydrophobic organic compounds from surface waters and is designed to collect suspended and dissolved phase organics.

2.1 Correlations Between Turbidity and Suspended Solids

This section describes techniques traditionally used to measure turbidity and suspended solids in waters, how the two parameters relate to each other and to various environmental impacts, and why one cannot be routinely substituted for the other. An additional literature review is presented in Attachment F-1. The term total suspended solids (TSS), sometimes referred to simply as suspended solids (SS), encompasses both inorganic solids such as clay, silt, and sand, and organic solids such as algae and detritus. It is a measure of the dry weight of suspended solids per unit volume of water, and is reported in milligrams of solids per liter (mg/L) determined by filtering a known volume of water through a filter of specified pore size (45 µm), and then drying and weighing the material retained on the filter.) USEPA Method 160.2 is often used for this 'TSS' measurement. Although popularly called suspended solids (the terminology used in this report), this method is more accurately called nonfilterable solids (or residue), because the size of separation (about 0.45 µm) is not the same as the boundary between suspended and dissolved solids, which varies among molecules but is generally around 0.1 μm. Another drawback of this method is that laboratories often run this method using an aliquot (100 mL) of the sample provided (typically a 250-mL sample bottle), with the associated possibility that some of the solids have adhered to or adsorbed to the surfaces of the container and therefore the reported result has a low bias relative to the 'true' value. The method used by USGS to measure suspended sediment, ASTM Method D3977-97, may be preferable.

Turbidity is an optical property of water that causes light to be scattered and absorbed rather than transmitted in straight lines through the sample. It is caused by the molecules of water itself, dissolved substances, and organic and inorganic suspended matter.

Turbidity measurements can be used as an operational aid in monitoring dredging and backfill placement operations as an adjunct to more costly and time-consuming suspended solids measurements in a laboratory. The primary reason for wanting to use turbidity measurements instead of suspended solids is that turbidity measurements are quick. Nephelometric turbidity readings can be done in a matter of minutes. On the other hand, taking a sample, transporting it to the laboratory, filtering it, drying it, weighing it, and calculating the suspended solids value can take from 3 to 24 hours. In the meantime, the suspended solids of the discharge or water body of interest will have changed. Therefore, laboratory measurements for suspended solids cannot be easily used to detect and correct short-term problems or performance standard violations. Because of this reason, turbidity measurements have historically been substituted for suspended solids. Turbidity is easy to measure quickly, but there is no universal relationship between it and suspended solids, nor among turbidity measurements made on different water-sediment suspensions, nor even among turbidity measurements made on the same suspension with different instruments. In addition, turbidity does not correlate well with many categories of environmental impact. However, turbidity can be used to indicate suspended solids concentration on a site-specific basis, if certain specific techniques are used.

Theoretical considerations prevent any simple, universal relationship between suspended solids and turbidity from ever being developed, because they measure different things, and their values are functions of different variables. Suspended solids depends on the total weight of particles in suspension, and is a direct function of number, size, and specific gravity of the particles, while turbidity is a direct function of the number, surface area, and refractive index of the particles, but is an inverse function of their size (for constant suspended solids) (Thackston and Palermo, 2000).

The problems in correlating turbidity and suspended solids are primarily due to two factors. First, the conversion from turbidity to suspended load involves a calibration that changes with changes in grain size of the sediment. Second, the calibration is changed as well by sediment color. A landmark paper co-authored by the inventor of one of the most widely used turbidity meters noted a factor of 10 change in calibration based on color alone, and an additional change in calibration that is linear with sediment grain size (Sutherland et al., 2000). For example, the calibration would change by a factor of 20 between white 5 micron sediment particles and gray 10 micron sediment particles. Such changes in sediment properties are not uncommon in nature. Since sediment color and grain size are not generally known during the course of a monitoring period, spot calibrations from samples are likely to contain unknown errors as sediment properties change in space and time (Agrawal and Pottsmith, 2000). These errors can reach several hundred percent, and greater. Laser sensors described below (under In-situ Total Suspended Sediment Measurement) overcome both these errors and advance the science of suspended sediment monitoring a quantum step forward.

2.2   In-situ Turbidity Measurement

Turbidity is the apparent "cloudiness" of water produced as light is scattered by particulate matter or dissolved material in the water. Presently established methods for measuring suspended sediments via optical turbidity are rooted in the pioneering work of Whipple and Jackson around

the year 1900 that lead to a candle-based turbidity standard called the Jackson Turbidity Unit (JTU). Devices commonly used to measure turbidity include the Jackson candle turbidimeter, absorptimeters, transmissometers, and nephelometers (McCarthy, Pyle, and Griffin 1974). All but the nephelometer measure the effects of both absorption and scattering of light. The nephelometer measures scattered light only, and is the most commonly used device in colloidal chemistry, drinking water treatment, and water quality management. Turbidity measured by such an instrument is reported in nephelometric turbidity units (NTU).

A transmissometer projects a narrow beam of light through a volume of water and measures the intensity of the beam as it exits the volume of water. If particles are in the water, they will attenuate the beam of light such that the light exiting the volume is less than the light entering the volume of water. The amount of attenuation can be measured, and with the appropriate calibration, these measurements can be used to estimate suspended-particle concentrations using empirically derived calibration curves. At low particle concentrations, transmissometers are very sensitive to small changes in particle concentration and/or size; however, at high-particle concentrations, transmissometers become saturated and lose their sensitivity to variations in concentration. Therefore, while transmissometers are very useful at measuring low-particle concentrations, they are inadequate for measurements at suspended solids levels above approximately 150 mg/L (Zaneveld, Spinrad, and Bartz 1979).

Nephelometers project a beam of light into a volume of water and measure the amount of light scattered out of the beam. The amount of light scattered is almost entirely dependent on the amount and size of particulate matter present in the volume of water. Ideally, a nephelometer would measure the amount of light scattered at all angles. Such a nephelometer is impractical, however, and standard nephelometers measure the scattered light at only one angle. Nephelometers use a device such as a photomultiplier tube or silicon photodiode to measure light that has been scattered at a specific angle, usually 90 degrees, from the main light path. The light source is usually a tungsten filament lamp or a light-emitting diode, and the light path is designed to minimize stray light falling on the detector. Thus, a zero signal means no light scattered at 90 degrees from the main light path and implies no turbidity.

Nephelometers used for in situ measurements are, in general, referred to as optical backscatter sensors (OBSs). OBSs measure the amount of infrared light backscattered from a volume of water. While suspended sediment will reflect infraenergy, organic matter will not (Tubman 1995). This characteristic of OBSs makes them well suited for measurement of sediment plumes because it does not bias the data by including organic matter. Since an OBS measures backscatter, its design is simple and compact relative to that of a transmissometer. More importantly, an OBS is capable of measuring much higher particle concentrations than a transmissometer, though it lacks the accuracy of the transmissometer at low-particle concentrations. Like the transmissometer, particle concentrations in the water can be estimated from OBS measurements using empirically determined calibration curves.

The ability of a particle to scatter light depends on the size, shape, and relative refractive index of the particle and on the wavelength of the light (Lillycrop, Howell, and White 1996). The reading on the instrument depends on many design parameters, including the light source, detector, electrical circuit, sample container, and optical arrangement. Therefore, two samples

with equal suspended solids concentrations but different size distributions of particles will produce very different turbidity readings on the same nephelometer; and two different nephelometers may produce different turbidity readings on the same sample, even if they were calibrated on the same standard (Vanous 1978; Hach 1972). Although the original Jackson candle turbidimeter was standardized with a specific fine silica suspension in which one JTU equaled 1.0 mg/L of suspended solids, modern turbidimeters are no longer standardized against the Jackson candle, and the term JTU is no longer used. The Jackson candle turbidimeter is no longer an accepted standard method (*Standard Methods* 20th edition, APHA et al.).

Modern turbidimeters are standardized against a formazin suspension with a value of 40 NTU. The standards should be prepared according to *Standard Methods* 20th edition (APHA et al.). The 400-NTU stock suspension should be prepared monthly, and the 40-NTU standard turbidity suspension should be prepared daily. Experience shows that this turbidity can be repeatedly prepared within an accuracy of ± 1 percent (Hach 1972). The formazin turbidity standard is assigned a value of 40 NTU and can be diluted to any desired value.

One of the main benefits of measuring turbidity is that turbidity sensors are relatively simple, inexpensive, and robust. The objective of most turbidity measurements is to identify the presence of suspended solids and quantify the suspended solids based on a correlation between turbidity and suspended solids. Historically, the standard practice has been to use turbidity measurements to estimate suspended solids. Such estimates are accurate only under the following conditions:

a. All measurements being compared are made with the same turbidity sensor.
b. The turbidity sensor is calibrated with a reference standard and suspended material from the area where the measurements are being taken.
c. Particle size and composition of the suspended material do not change significantly during the measurement period.

Turbidity can also be measured in the field by collecting water samples and using portable instruments to analyze the samples. While these instruments are typically less expensive than in situ sensors, the measurements take longer and may not represent true in situ conditions since particles may settle out of suspension prior to analysis. The cost of these instruments is approximately $1,500 to $2,000.

2.3 In-situ Total Suspended Sediment Measurement

Historically, suspended solids has been measured by collecting water samples and analyzing these samples in an offsite laboratory. Water samples can be collected using a bottle sampler or a submerged pump. Independent of the collection method, care must be taken to ensure that suspended particulate matter does not settle out of suspension or flocculate during collection or prior to analysis. Offsite laboratory analysis is time-consuming and cannot provide data in the short term. However, this approach is considered to be the most accurate and reliable method for measuring suspended solids. The alternative has been to estimate suspended solids based on other measurements such as turbidity or acoustic backscatter, both of which have limitations as discussed above.

Recently, instrumentation has been developed that provides an alternative for measuring suspended solids in situ more accurately than can be achieved by using correlations with turbidity. Laser In Situ Scattering and Transmissometry (LISST) measures the scattering of a laser beam by particles in a volume of water. It should be noted, that laser diffraction measurements have been used to measure and characterize suspended sediments and floc-sizes in situ since 1985 (see, e.g. Bale and Morris, 1987; McCabe et al.,1993; Agrawal and Pottsmith, 1994; Gentien et al.,1995; Bale, 1996; van der Lee, 1998).

The LISST-25 is a small, self-contained unit suitable for field deployment with real-time data return capabilities (Sequoia Scientific, Inc.). The instrument is capable of measuring particle total volume, particle total area, and Sauter mean diameter within a particle range of 1.2 to 250 mm. These parameters are defined as follows:

a. Particle total volume is the volume of material per volume of water.
b. Particle total area is the projected cross-sectional area of the particles per volume of water.
c. Sauter mean diameter is the ratio of the particle total volume to the particle total area.

If the density of the suspended particulate matter is assumed, calculating suspended solids by multiplying the particle total volume by the assumed density is possible. Other models in the product line of the LISST instrument are also capable of measuring the particle size distribution. The LISST-100 is the first in situ laser that simultaneously measures the beam attenuation coefficient, the volumetric concentration (ml/L) and in situ particle size spectra. It is designed to be submerged to a depth of maximum 300 m and equipped with a built-in datalogger.

The LISST-100 measures the particle size distribution in 32 logarithmically spaced size classes in the range 1.25 to 250 μm (a LISST-100 type B). Other versions of the instrument can measure size ranges of 2.5 to 500 μm and 7.5 to 1,500 μm, in all cases spanning a 200:1 dynamic range. A detailed description of the design and the operational principles of the LISST-100 can be found in Agrawal and Pottsmith (1994), Agrawal et al.(1996) or Traykovski et al. (1999). However, the basic principles are explained very briefly below. The LISST-100 measures the angular distribution of forward scattered light energy over a path length of 5 cm, using a collimated laser beam with a wavelength of 670 nm. The energy of the scattered light is detected on 32 logarithmically spaced ring detectors and stored in a built-in datalogger. When data collection is complete, these raw data are offloaded and mathematically inverted. The inversion yields the area distribution of the suspended particles (in 32 size classes). By multiplying the area distribution by the diameter of each size class, the particle volume distribution is obtained. Summing the volume distributions in all 32 size classes and dividing by an instrument-dependent calibration constant, the absolute volume concentration (ml/L), is found. The part of the light not scattered is detected by a photo-diode in the center of the ring detector, thus yielding the optical transmission, T, of the water. From the optical transmission the beam attenuation coefficient at 670 nm, c(670), can be calculated using Eq. (1)

$$c(670)(m^{-1}) = -1/0.05 \text{ m } \times \text{ ln } (T) \dots\dots\dots\dots\dots\dots\dots\dots\dots\dots(1)$$

The processed data output from the LISST-100 thus consists of a particle volume distribution (in 32 size classes), an absolute volume concentration (ml/L) and a beam attenuation coefficient at

670 nm. Furthermore, the LISST-100 records the temperature and pressure. From the particle volume distribution, statistical parameters such as the mean and standard deviation can then be calculated. All software necessary for obtaining and analyzing raw data is supplied by the manufacturer of the LISST-100, Sequoia Scientific Inc., USA. (See Figure 1.)

Although the LISST instruments have not been used extensively in field studies of plumes when compared with turbidimeters, some very good documented information on their performance does exist (Melis, T., 2002, Mikkelsen, O., 2000). A recent study comparing the LISST to traditional methods of measuring suspended-sediment concentration found the LISST provided accurate measurements of total volume concentration of suspended sediments (Traykovski et al, 1999). Once the accuracy and limitations of these systems have been thoroughly documented by site-specific testing at the Hudson River PCBs Superfund Site during the two to three years baseline/pre-dredge monitoring period, this instrument could prove very useful for in situ monitoring of sediment plumes in the Hudson River during Phase 1 and Phase 2. The cost of the monitoring equipment is approximately $15,000 to $30,000 for the LISST-25 and LISST-100. Because of the cost some limited use of these instruments is warranted such at the far-field stations and for daily readings at the near-field stations.

2.4 Semipermeable Membrane Devices (SPMDs)

An SPMD is a passive sampling device that consists of a thin film of the neutral lipid, triolein, sealed inside a layflat, thin-walled tube of nonporous *(i.e*, no fixed pores; only transient thermally mediated cavities) low-density polyethylene (LDPE). The diameters of the transient cavities range up to approximately 10 Å, effectively precluding sampling of any contaminant molecules associated with dissolved organic matter or particulates. This cavity size limitation has an important consequence: in general, only dissolved chemicals with molecular masses less than about 600 are sampled by SPMDs and this molecular mass limitation is very similar to that imposed by the pores of biomembranes.

At saturation, the capacity of the SPMD for a hydrophobic compound (like PCB) is generally related to the compound's octanol-water partition coefficient (K_{ow}). The higher a compound's K_{ow}, the greater the capacity of the SPMD for that compound. Due to the very high concentration factors attained, even ultra-trace levels of the hydrophobic contaminants are readily analyzed. Standard SPMDs are designed to sequester and concentrate hydrophobic compounds like PCBs and PAHs. SPMDs are not designed to concentrate ionic species such as ionic metals, salts of organic acids, or very polar organic chemicals. Neutral organic chemicals that are hydrophobic (i.e., with log K_{ow} values $\geq$ 3) will be concentrated significantly above ambient levels. In reality, any compound with a log $K_{ow} \geq 1$ will be concentrated by the SPMD, but for compounds with log K_{ow} values ≤ 3, there is no significant advantage in using SPMDs in preference to other sampling techniques.

When placed in an aquatic environment, SPMDs passively accumulate hydrophobic organic compounds, such as PCBs. The LDPE tubing mimics a biological membrane by allowing selective diffusion of organic compounds. Triolein is a major nonpolar lipid found in aquatic organisms. The passive sampling of the hydrophobic organic chemicals is driven by membrane- and lipid-water partitioning. (See Figure 2.)

SPMDs can be deployed for long periods of time (days to months) and used to estimate the time-weighted mean concentrations of the hydrophobic organic compounds in the water body. The SPMD is placed on a rack, which is inserted within a protective "shroud," and is then ready for use in the water. An SPMD can also be used vertically and horizontally as illustrated below:

An SPMD will effectively sample 0.5 to 10 L per day, depending on the chemical's hydrophobicity (as quantified by its water solubility or octanol-water partitioning coefficient, K_{ow}) and other factors. A compound with log K_{ow} 6 would need 200 days at an effective sampling rate of 10 L per day to reach 90 percent of equilibrium. However, during the first 50 days, the uptake rate into the SPMD is linear.

The concentrations of these chemicals in rivers can change daily or even hourly. To get a true picture of the amount of contaminants present in the water column, many samples would have to be collected and analyzed. The SPMD allows the calculation of a cumulative time-average of the concentration of each contaminant while the SPMD was in the water.

The ambient "truly dissolved" water concentration (C_w) can be estimated based on the concentration in the SPMD (C_{SPMD}), the volume of the SPMD (V_{SPMD}), the effective sampling rate (R_s), and the time of deployment (t):

$$C_w = C_{SPMD} \, V_{SPMD}/ \, (R_s * t)$$

After a typical deployment period of approximately 15 to 30 days, the SPMDs are removed from the aquatic environment and recovered by dialysis with a nonpolar solvent such as hexane. This extract is then reduced, cleaned up, and enriched. The cleanup procedure typically includes gel permeation chromatography. This process removes any lipid and polyethylene waxes that might have carried over during the dialysis extraction. Further cleanup can be effected during enrichment on an activated alumina and silica gel column. The enriched extract is then analyzed for target compounds using chromatographic techniques.

A major portion of the sequestered residues can be recovered by opening the ends of the SPMD polyethylene tube and rinsing out the lipid with an organic solvent. However, analytes are generally recovered by dialyzing the intact SPMD (which requires removing periphytic growths, minerals, and debris from the exterior membrane surface) in an organic solvent such as hexane. Using this approach, contaminant residues present in the membrane (sometimes representing as much as 50 percent of the total) are also recovered for analysis and the dialysis process separates nearly all of the bulk lipid from the chemicals of interest.

One of the problems encountered with deployment is biofouling, the coating found on the membrane exterior. This biofouling layer can impede flux across the membrane, thus slowing the effective sampling rate (R_s). This impedance factor is specific to each SPMD at any given point in time. Impedance for a specific deployment can be quantified by measuring the loss of a surrogate compound (contained within the SPMD) during deployment.

The SPMD sampling rates are directly proportional to the SPMD membrane surface area. For example, a standard 1-g triolein SPMD (surface area about 450 cm^2) may extract 5 L of water

per day for a PCB congener, whereas a standard triolein SPMD with half the surface area (225 cm^2) (0.5-g of lipid) can be expected to extract 2.5 L of water per day of the same congener, assuming similar conditions for the exposures.

Due to the highly sensitive nature of the SPMDs, assembly and placement of the devices requires considerable care. According to Huckins *et al.* (1996), the following quality control (QC) procedures must be followed during the SPMD preparation phase:

$\quad$ Synthetic triolein or lipid are used, and all new lots or batches are analyzed for contaminants, ampulated and stored in a freezer until use;

$\quad$ Accurate delivery of small volumes of triolein requires the use of a micropipettor equipped with a total displacement plunger;

$\quad$ SPMD tubing is batch extracted with nanograde hexane or cyclohexane, just prior to use in SPMD construction;

$\quad$ Enclosure of triolein in SPMD layflat tubing is achieved using a heat sealer which results in a molecular weld;

$\quad$ After assembly, SPMDs are sealed in clean, gas-tight paint cans (solvent rinsed to remove cutting oils) or gas phase sampling bags (Tedlar®) for transport to deployment sites.

Placement of the devices is important because of a variety of factors. According to Huckins *et al.* (1996), the following quality control (QC) procedures must be followed during the deployment phase:

$\quad$ Use of plastic components should be minimized, except for Teflon, due to the possible presence of leachable organic residues;

$\quad$ The design of the structure to hold the SPMD should minimize abrasion of the membrane; and

$\quad$ Since the SPMD membrane generally controls uptake, current velocity is usually only a concern in terms of abrasion and tethering.

Another important phase to consider is the recovery and storage of SPMDs. According to Huckins *et al.* (1996), the following QC procedures must be followed during this phase:

$\quad$ As soon as SPMDs are recovered from the environment, they should be sealed in the original can or Tedlar bag and placed on ice. The devices should be shipped to the processing laboratory overnight; and

$\quad$ SPMDs should be stored in the original container at -20° C until they are analyzed.

During the dredging in the Hudson River, SPMDs will be deployed at the far-field stations for periods of 15 days. The dissolved phase PCB concentration in the water column over the two-week period can then be determined. It should be noted that these measurements should be

regarded as qualitative and used to measure relative changes in the water column concentration over successive two-week periods.

2.5   Trace Organic Platform Sampler (TOPS)

The detection of trace organic compounds in the water column is generally very problematic because many target compounds are typically present at concentrations that are below the detection limits of conventional analytical methods. In these instances, a non-detect result generally represents a failure in field sampling and/or laboratory analysis to measure these target compounds at environmentally relevant concentrations. Available analytical methods require large volume samples to resolve concentrations in the picogram to femtogram per liter range. In environmental settings where concentrations are known to be exceedingly low, collection of large grab samples can be logistically difficult and cumbersome. Field processing of samples in these settings greatly simplifies the collection process while significantly lowering detection limits.

In order to overcome these difficulties, the New York State Department of Environmental Conservation (NYSDEC) developed the Trace Organic Platform Sampler (TOPS) as a tool to obtain water column samples. TOPS is a set of plumbing, pumps, and sensors that concentrates hydrophobic organic compounds from surface waters. The TOPS is designed to collect suspended and dissolved phase organics. The TOPS uses glass fiber cartridge filters (1 micron pore size) to capture suspended solids and the synthetic resin Amberlite XAD-2 (XAD) to capture dissolved phase hydrophobic organic compounds like PCBs. The 1 micron pore size filters were chosen because they are readily available in desirable configurations and they were assumed to be efficient at capturing most of the suspended solids in river settings.

XAD is a polymeric adsorbent of hydrophobic cross-linked polystyrene copolymer supplied as 20-60 mesh beads. The beads are an agglomeration of many microspheres giving a continuous gel phase and a continuous pore phase. The XAD surface area is 300 m^2/g. The open cell porous structure allows water to easily penetrate the pores of the resin. In the adsorption process, the hydrophobic portion of the adsorbate molecule is preferentially adsorbed on the hydrophobic polystyrene surface of the resin while the hydrophilic section of the adsorbate remains oriented in the aqueous phase. Compounds adsorbed do not penetrate into the microsphere phase and remain at the surface where they can be easily eluted. Unlike liquid/liquid extraction procedures, it is easy to scale up XAD sampling systems to treat exceptionally large volumes of water. These large water volumes have a greater likelihood of containing a detectable mass of target organic analyte than smaller volumes.

The best use of TOPS is for obtaining whole water concentrations of extremely dilute hydrophobic organic compounds.. With adequate support, TOPS is a very powerful field tool that can be deployed from ships or fixed locations where sample size is unlimited. In such cases, there is virtually no detection limit as more analyte can be obtained simply by pumping more water. TOPS typically processes more than 5,000 liters in order to achieve adequate detection of target compounds. Where field setup is inconvenient and concentrations are expected to be relatively higher, TOPS can be used in bench-top mode. Samples on the order of tens of liters can be brought in from the field and batch processed.

In its original configuration, the TOPS was run by an on-site operator for a fixed length of time (as short as one day) or at fixed intervals to sample wastewater effluent, coastal waters, and other low suspended sediment environments. USGS, in cooperation with NYSDEC, modified the TOPS for operation in river environments where suspended sediment concentrations are relatively high. Additional TOPS modifications allow for remote, automated, and flow-weighted operation (USGS, 2003 and NYSDEC, 2003).

The TOPS uses 110 VAC and processes water through cartridge filters (available in 4 or 10 inch lengths), and through XAD columns at a maximum rate of 620 mL/min. TOPS can process water at a much greater rate through the filter (3,200 mL/min) than through the XAD so significant amounts of suspended solids may be captured even in waters with low suspended solids. Since the pump rates through the glass fiber filters and through the XAD are independent, sampling rates can be adjusted depending on the turbidity of the water.

Remote and automated operation was made possible by adding a Campbell CR10X datalogger that monitors stream stage, triggers sample collection based on stream discharge, and monitors flow through the XAD resin and filter as well as backpressure associated with the filter. A modem connected to the datalogger allows a user to dial into the site to initiate, monitor, or stop sampling. Hydrologic events rarely occur at convenient times so datalogger programming includes a set of conditions under which TOPS sampling will start automatically. These conditions usually take the form of a threshold change in river stage over time, but could include a variety of other programmable triggers including river discharge. As with starting a TOPS sample, stopping can be accomplished either manually or automatically. Automatic termination based on river stage is set for when the stage falls 80 percent of the difference between the event start stage and peak stage.

Collection of composite samples during periods of changing river discharge is best accomplished by flow-weighting the volume of water collected. Flow-weighting is a method by which the volume of sample water collected is proportional to the volume of water passing the sample station. Flow-weighting, as compared to fixed interval sampling, avoids over-representing conditions present during the beginning and end of the event, and likewise under-sampling the mass flux of contaminants passing during the hydrologic peak. Contaminant concentrations derived from flow weighted sample collection are easily used for determining contaminant flux by multiplying them by the mean river discharge during the period of sampling and converting to the appropriate units. In practice, flow-weighting is accomplished by collecting a fixed volume or sub-sample of river water every time a pre-set volume of river water passes the sampling station. This pre-set river water volume is an educated guess based on the anticipated river discharge maximum, expected duration of the event, and minimum sample volume required. Real time discharge data is required to collect a flow-weighted sample. The interval that discharge data is collected is dependent upon a variety of factors, but is principally dictated by the pre-set volume of river water used to trigger a sub-sample; in NYSDEC's application under the Contaminant Assessment and Reduction Project (CARP) discharge data were typically collected once per minute.

To allow sampling when suspended sediment concentrations are high, another pump was added to the sampling system that delivers a flow-weighted sample to a settling/compositing tank. In this configuration, the TOPS draws water from the tank instead of pulling water directly from the river. The tank sits on a scale which is monitored by the datalogger. The mass of the water in the tank is used to control when the TOPS turns on and off and when the river water pump should turn off. The tank allows material that would otherwise clog the TOPS filter prematurely to settle. Settled material in the tank is collected and filtered at the end of the event and composited for analysis with the TOPS filter. The advantages of using an additional pump in the sampling process include: (1) the use of pumping rates that keep material in suspension without compromising the integrity of the TOPS filter; (2) the ability to purge the sampling line before and after a sampling interval; and, (3) the removal of the TOPS from the role of collecting a flow-weighted sample.

The addition of the tank to the TOPS sampling system is primarily designed to extend the life of the TOPS cartridge filter - material settling to the bottom of the tank avoids TOPS filtration thereby reducing the amount of material on the filter and prolonging filter life. Besides this obvious advantage, the tank has several additional benefits that improve the quality of sample collection. Without the tank, the main TOPS pump must collect and process the sample directly from the river, this requires the main pump to pull water from the river at a rate of at least 2 ft/sec to keep material in suspension. The filter may be able to process the volume of water required, but when the filtration is time constrained, the result is an increase in backpressure from the filter to the point where the TOPS shuts down. Additionally, as the filter accumulates sediment and backpressure builds, the effective pumping rate decreases with time – this introduces bias into the sample collection in that the efficiency of the point intake to collect suspended material changes over time.

By removing the main TOPS pump from serving as the direct collector of river water, the pump rate of the main TOPS pump can be significantly slowed. Slower filtration itself reduces backpressure from the cartridge filter thereby extending processing time. Slower pump rates also reduce the formation of air bubbles in the sampling line produced from degassing of sample water under rapidly changing pressure conditions. Air bubbles can adversely impact the accuracy of the flow meters, which are critical in determining contaminant concentrations. The tank also buys the operator time to get to the site in the event that maintenance is needed. Sub-samples can be composited in the tank at the beginning of the event before and during installation of the TOPS cartridge filter and XAD and during the event to change a clogged filter. By remotely monitoring river conditions and TOPS backpressure, sub-samples can be collected without interruption over the course of the hydrologic event.

A further advantage the tank and sub-sample pump combination have over direct TOPS pumping is that the sub-sample pump can flush excess sub-sample water remaining in the line following collection of a sub-sample without adversely affecting TOPS processing or pumping sample water back to the river. Without intake line flushing, the sub-sample collected directly by the TOPS may be partially or entirely made up of water that remains in the sample line from the previous sub-sample. In addition, part or all of the sample water collected may not adequately represent the suspended sample fraction in that settling of suspended material occurs in the

sample intake line between sub-samples – this is particularly a problem in locations requiring long sections of vertical or near vertical sample line.

Wound glass fiber cartridge filters are capable of filtering large volumes of water without clogging, but have the disadvantage of allowing more suspended material to pass relative to conventional plate filters with the same nominal pore size. Experiments conducted to test the efficiency of the 10 inch cartridge filters (both 0.5 and 1 micron nominal pore size) indicate the efficiency changes with the volume of water processed, often times in unexpected ways, but generally in response to material loading of the filter. Over the course of these tests, both filter pore sizes trapped between 85 and 89 percent of the total mass of sediment sampled with pre-filter concentrations ranging from 3 to 82 mg/L. The TOPS can be equipped with a series of solenoid valves to periodically divert a sub-sample of water to a sample container. These valves and containers can be placed after the filter to assess the overall trapping efficiency of the filter.

A conventional automatic sampler is used with the TOPS to help interpret and support the organics data collected by the TOPS. This sampler collects discrete sample pairs for analysis of suspended sediment concentration and particulate and dissolved organic carbon concentration. Sediment and organic carbon samples are collected at the beginning, end, and peak of the hydrologic event in addition to measured changes in stage (e.g. every 0.5 feet of stage change).

2.6 ISCO Portable Water And Wastewater Sampler

All the portable samplers manufactured by ISCO can be divided into two groups, the full-size sampler and the compacted sampler. The compacted samplers are specially designed for the locations with limited access such as manhole. The full-size sampler needs a larger space for installation. The open-channel flow conditions at the far-field monitoring stations in the Hudson River would require the fitness of the full-size sampler.

The 3710 composite-only portable sampler combines simple operation and high volume capacity for single-bottle sampling. It collects composites samples – based on time or flow interval – in a 2.5 gallon glass or polyethylene bottle or a 4 gallon polyethylene bottle. Up to 24 sampling stop and resume times can be preset for unattended, automatic sampling. The controller can be easy setup for uniform time interval, non-uniform time interval and flow-paced sampling with or without time delay.

The full-feature 3700 Sampler collects sequential or composite samples based on time, flow rate, or storm conditions. It is an ideal choice if the parameter monitoring and logging capabilities are not needed. The exclusive LD90 gives the automatic compensation for changes in head height, plus automatic suction line rinsing to prevent cross contamination. Basic and extended programming modes are provided for uniform time intervals, non-uniform time intervals, stormwater runoff sampling, multiple bottle compositing and split sampling. The bottle configurations for composite sampling are the same as for the 3710 Sampler. Sequential sampling bottle configurations include 24 x 1 liter polypropylene or 350 ml glass, 12 x 1 liter polyethylene or glass and 4 x 1 gallon polyethylene or glass.

Both the 3710 and 3700 pumps maintain the USEPA-recommended 2 feet per second line velocity at head heights up to 16 ft, with ¼-inch suction line. For higher lifts, the 6700 series is recommended. The 6712 Portable Sampler is the most sophisticated full-size sampler. Samples can be delivered at the USEPA-recommended velocity of 2 feet per second, even at a head height of 26 feet. The plug-in 700 Series Modules and the new SDI-12 interface make it easy to add flow and parameter monitoring to the basic system. The advanced 6712 Controller allows the user to select different programming modes to assure the most suitable routine for specific application. Standard 4MB of memory gives the user great flexibility for logging environmental data. Choice of 11 different glass and plastic bottle configurations ranges from 24 x 1 liter to 1 x 5.5 gallon.

All the samplers require the power of 12 VDC. Ni-cad lead-acid batteries can be purchased from ISCO. But depending on the sampling frequency and the volume of one sample, the battery can last only 1 to 3 days. To meet the 2-weeks continuous sampling requirement as set for the routine monitoring, tying the electricity to the sampling location to provide the power for the sampler will be the most convenient and economic way given by the duration of the project and the number of samples to be collected. The purchasing cost is $1975 for Model 3710, $2425 for Model 3700, and $2700 for Model 6712. To analyze PCB appropriately, the laboratory requires a 16-L sample. The 5-gallon container is needed to collect sufficient amount of water sample. Given the features of these samplers and the needs of this project, Model 3700 and Model 3712 would be the better choice. The details regarding how to deploy the samplers during remediation monitoring should be fully addressed in the design phase.

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Figures
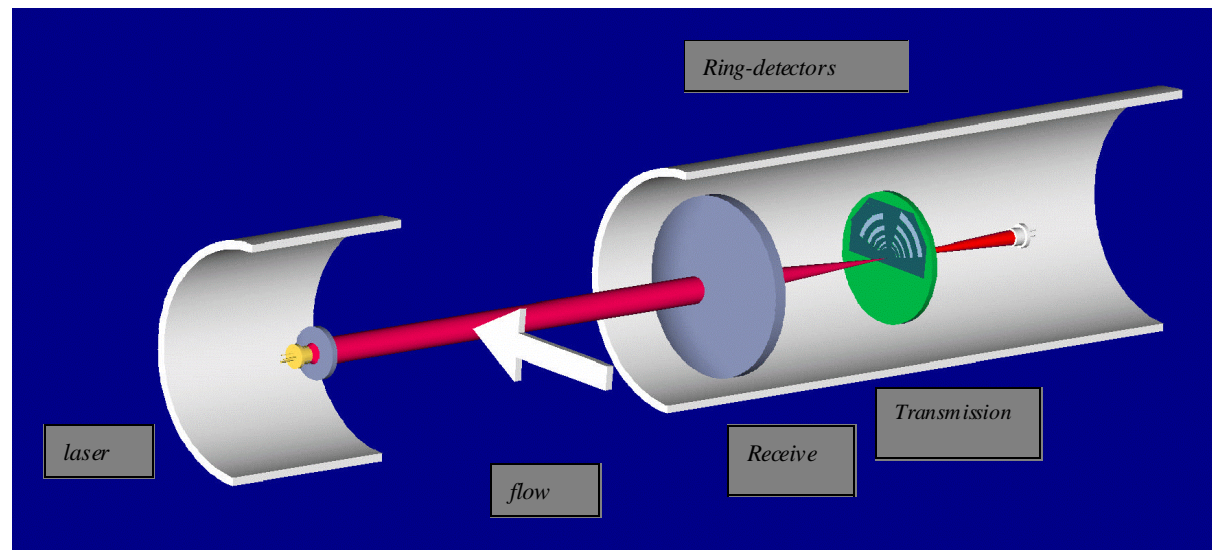
Figure 1 Laser diffraction principles – a cut away view of the basic LISST-100 instrument.

A collimated laser beam illuminates particles (left to right). Multi-angle scattering is sensed by a specially constructed photo-diode array placed in the focal plane of the receiving lens. The array detector has 32 concentric rings, placed in alternate quadrants. An aperture in the center passes the attenuated beam for measurement of optical transmission.
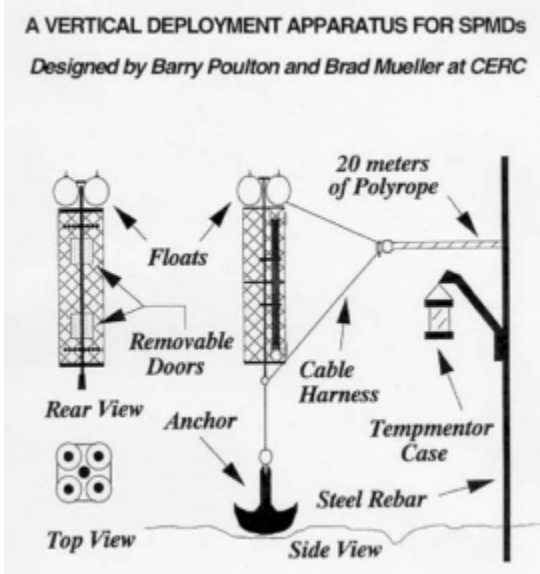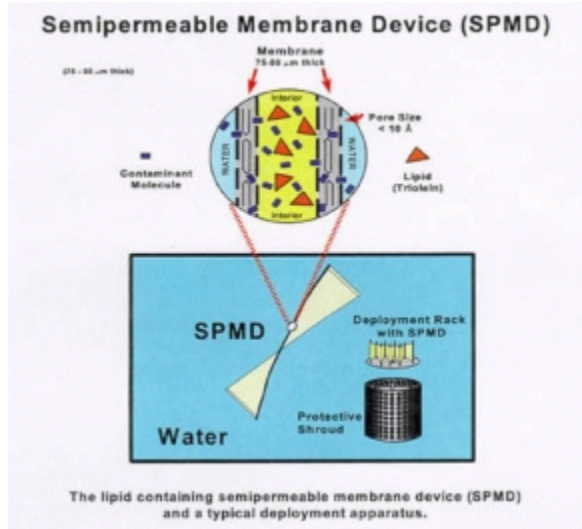
Figure 2. SPMD Apparatus

Attachment F-1

Literature Review

Attachment F-1
Literature Review

1. Introduction

PCB concentrations cannot be measured quickly or easily in the field, requiring time-consuming laboratory analyses. Turbidity and total suspended solids (TSS) can be determined relatively quickly and easily using real-time monitoring devices. To develop an estimate of real time PCB concentration in the vicinity of the dredging operations, the development of relationships between turbidity and TSS and TSS and PCB concentrations will be investigated.

Analysis of TSS and PCB data from a set of GE water column monitoring samples did not yield a correlation between the two parameters. Based on this observation, the PCB concentrations in the near-field will be projected using modeled solids concentrations (obtained using the DREDGE and/or SED20 models), consideration of the travel time, average concentrations in each river section, and an estimate of the time to reach equilibrium between the dissolved and suspended phases. It is not anticipated that PCB concentrations will be measured in the near-field during remediation.

PCB concentrations will be measured at the far-field stations, via sampling and analysis, and the levels will be compared with the TSS levels from the near-field stations to determine if a correlation exists. Phase 1 of the remediation will provide information that can be used to further refine any observed relationship between near-field solids and far-field PCB concentrations; refinements could be incorporated in the Final Phase 2 Engineering Performance Standards. The papers below were reviewed to investigate the feasibility and applicability of such a correlation.

2. Paper List

1. Chattooga River Watershed Ecological/Sedimentation Project (Pruitt at al., 2001)

2. Improved Methods for Correlating Turbidity and Suspended Solids for Monitoring (Thackston et al., 2000)

3. St. Lawrence River Sediment Removal Project Environmental Monitoring Plan: Section 2: Pre-Sediment Removal Data Collection (BBL Environmental Services, Inc., 1995)

4. Use of Acoustic Instruments for Estimating Total Suspended Solids Concentrations in Streams—The South Florida Experience (Patino et al., USGS, 2003)

5. Appendix K: Water Quality Monitoring Pre-Design Field Test Dredge Technology Evaluation Report, New Bedford Harbor Superfund Site, Section K.6.2 (USACE, 2001)

6. Suspended Solids Flux Between Salt Marsh and Adjacent Bay: A Long-term Continuous Measurement (Suk et al., 1999)

3. Chattooga River Watershed Ecological/Sedimentation Project (Pruitt at al., 2001)

The purpose of this study was to conduct a sediment yield evaluation and analyses to determine if sediment was a primary cause of physical and biological impairment to streams within the Chattooga River watershed, located in northeast Georgia, northwest South Carolina, and southwest North Carolina. This was done by sampling sediments and aquatic ecology from different areas of the watershed and correlating the data by site.

For the aquatic ecological analysis, a total of three reference sites and 56 other sites from six subwatersheds (Headwaters, Lower Chattooga, Middle Chattooga, Stekoa Creek, West Fork, and Warwoman Creek) were sampled. Biological sampling methods were focused on benthic macroinvertebrates and used modified rapid bioassessment protocols. Reference sites were chosen prior to sampling based on habitat condition, *in situ* water chemistry, and surrounding land use; two sites were located on the Chattooga River and one on the upper Chattahoochee River located outside of the Chattooga watershed. Data from all stations were analyzed using a multimetric approach; 17 metrics were calculated from the raw data, and ultimately the five of those that had the greatest ability to detect impairment were selected.

For sediment sampling, 17 stream reaches were selected for storm flow investigations based on the following criteria: relative degree of biological impairment as measured using modified rapid bioassessment protocols, position within the watershed, relative geomorphic condition, and access logistics. Storm flow investigations were performed during three storm events in March 1998, June 1999, and March 2000. A total of 58 observations were made across the 17 stations.

Total suspended sediment (TSS) was analyzed through the filtration of whole water samples and in accordance with USEPA Method 160.2. Bedload samples were collected using a 6-inch cable-suspended bedload sampler or a 6-inch wading type bedload sampler. The samples were transported to the laboratory in 1-liter containers, and processed for particle size determination in the laboratory using the EPA-SESD wet sieve method. Laboratory results of dry-weight bedload samples ($M_{b, grams}$) were converted to bedload transport rate (Q_b, tons/day) by the following equation:

$$Q_B = K(W_T/T)M_T$$

where:
- Q_B = bedload discharge (tons/day)
- K = converts grams/second/foot to tons/day/foot
- W_T = wetted surface (ft)
- T = total time sampler on bottom (seconds)
- M_T = total mass of samples (grams)

The amount of bedload sediment measured over the course of the three storm events averaged 13.32 tons/day, with mean particle sizes ranging from fine sand to very coarse sand. On average, the bedload sediments only accounted for 14% of the total sediment load. The TSS averaged 85.3 tons/day over the course of the three storm events, making up 86% of the total sediment load on average. Total sediment load (bedload sediment + TSS) was compared to discharge and

road density (road length/corresponding drainage area). Road density is a factor that represents the net impacts of road construction and maintenance, interception of subsurface interflow, routing of other non-point sources to the stream, and entrainment, mobilization, and transport of sediment to the stream.

Study results indicated that the biological conditions in most of the streams sampled showed little or no impairment due to sedimentation effects. 78% rated "very good" or "good," 19% rated "fair," and 3% rated "poor." None rated as very poor. Although some sedimentation or habitat effects of sedimentation were evident at many sites, a negative biological response was not always presented. The most degraded biological community was observed in the Stekoa Creek subwatershed. Data indicated that impaired streams contained a higher concentration of bedload and suspended load sediments when compared to the reference streams. Study results also indicated that the road density and sediment sources associated with the road density were source of 51% of the total sediment loading.

Good correlation was observed between the biological index and the normalized TSS data. Data suggest that a TSS concentration normalized to discharge/mean discharge greater than 284 mg/l adversely affected the biological community structure. However, based on regional concentrations, a normalized TSS concentration of 58 mg/l or less during storm flow provides an adequate margin of safety and is protective of aquatic macroinvertebrates in the area. Corresponding turbidity limits of 22 and 69 NTU represent the margin of safety and threshold of biological impairment.

Reference

Pruitt, B. A.; Melgaard, D. L; Howard, H.; Flexner, M. C.; Able, A. "Chattooga River Watershed Ecological/Sedimentation Project," *FISC Proceedings,* Federal Interagency Sedimentation Conference, Reno, Nevada, March 26-30, 2001.

4. Improved Methods for Correlating Turbidity and Suspended Solids for Monitoring (Thackston et al., 2000)

This article describes techniques that are traditionally used to measure turbidity and suspended solids in water, how the two parameters relate to one another and to various environmental impacts, and why one cannot be routinely substituted for the other. This paper also outlines techniques describing the use of quick turbidity measurements as aid to monitoring dredging and dredged material disposal operations.

Turbidity and suspended solids are common parameters of concern for regulatory agencies, and thus are often included in the environmental monitoring plans for dredging operations. Because suspended solids measurements cannot be made quickly and easily in the field, turbidity measurements are often taken instead. While turbidity can be measured quickly, there is no universal correlation between the two parameters, or between turbidity measurements taken from different suspensions or the same suspension with a different instrument. However, turbidity can be used as an indicator on a site-specific basis.

Total suspended solids (TSS) include both inorganic solids and organic solids. TSS is a measure of the dry weight of suspended solids per unit volume of water, and is reported in milligrams of solids per liter of water (mg/l).

Turbidity is an optical property of water that causes light to be scattered and absorbed rather than transmitted in straight lines through the sample, and is reported in Nephelomatic Turbidity Units (NTUs). The source of turbidity in a sample includes suspended inorganic and organic matter, water molecules, and dissolved substances. The ability of a particle to scatter light depends on the size, shape, relative refractive index of the particle, and the wavelength of the light.

There is no universal correlation of TSS and turbidity, but sediment-specific correlations are useful as a real-time indicator of suspended solids. Such correlations have been developed in the laboratory using whole sediment samples. Generally, any samples used to produce a correlation between TSS and turbidity must be suspension-specific, not just site-specific. The sample must approximate the suspension to be representative of the size, number, shape, and type of particles present.

Most discharge or monitoring permits that are associated with dredging operations are based on TSS rather than turbidity because TSS correlates well with environmental impact and is at least roughly comparable from site to site and sediment to sediment.

It has been suggested that there are three general situations where a TSS-turbidity correlation curve may serve as an aid in the routine monitoring of a dredging operation:

- Solids resuspension in the immediate vicinity of the dredge (20-50m) where most solids will be continuously replenished by dredging actions.

- Containment area effluent, where only the finer particles will be present due to the settling of larger, heavier particles near the point of inflow for the contaminant disposal facility. For this case, a laboratory settling column and test procedure would be required to obtain a representative sample.

- Open-water dredged material placement, a case in which the larger, heavier solids will begin to settle to the bottom immediately upon leaving the dredge discharge pipe, hopper, or barge usually in a well-defined plume. This case requires the use of a laboratory column-settling test to obtain a representative sample.

Reference

Thackston, E. L.; Palermo, M. R. "Improved Methods for Correlating Turbidity and Suspended Solids for Monitoring," *DOER Technical Notes Collection* (ERDC TN-DOER-E8), U.S. Army Engineer Research and Development Center, Vicksburg, MS, 2000.

5. St. Lawrence River Sediment Removal Project Environmental Monitoring Plan: Section 2: Pre-Sediment Removal Data Collection (BBL Environmental Services, Inc., 1995)

The goal of the pre-sediment removal data collection program was to verify bottom conditions, obtain background water quality information, and obtain a location survey of the sediment control system in the St. Lawrence River at the GM Massena site. One of the tasks planned to accomplish these objectives was pre-dredging turbidity monitoring.

To perform real-time monitoring that allowed for a rapid response to changing river conditions, a water quality parameter that is easily measured and correlates with sediment resuspension during removal activities must be chosen. Turbidity was the parameter selected in this case.

A downstream total suspended solids (TSS) maximum limit of 25 mg/l above background was defined as the conservative action limit based on two variables: previous environmental dredging projects and a 1994 site-specific bench-scale laboratory correlation between TSS and turbidity.

The 1994 bench scale experiment established a site-specific correlation between TSS and turbidity for the GM Massena site, resulting in the use of real-time turbidity measurements as a surrogate for TSS measurements. The laboratory-produced correlation, which is based on a combination of all data points from the treatability test (including some elevated TSS results ($>$ 300 mg/l) from the beginning of the settling test), is described by the equation 1 below:

$$\text{Turbidity (NTU)} = 7.3745 + (0.61058 \text{ X TSS}) + (0.00094375 \text{ X TSS}^2) \quad (1)$$

with a correlation coefficient of $r^2 = 0.941$

Turbidity monitoring data collected in 1994 indicated that the St. Lawrence River can be characterized as having a relatively low suspended solids content (based on the evaluation of background river water samples, which contain $<$ 10 mg/l TSS) and low turbidity readings. A regression analysis was rerun by BB&L only including data that fell within the expected working range, defined as: TSS $<$ 60 mg/l and turbidity $>$ 60 NTU. The regression equation 2 calculated is defined below:

$$\text{TSS (mg/l)} = [0.63x \text{ x (turbidity in NTU)}] + 6.8 \quad (2)$$

with a correlation coefficient of $r^2 = 0.43$

Based on the revised regression (2), a turbidity of 28 NTU would correlate to a value less than 25 mg/l TSS concentration. Dredging activities would not take place when the measured TSS background was above 60 mg/l. So, due to the nearly linear relationship that exists between turbidity and TSS for the St. Lawrence River in the subject area, a turbidity increase of 28 NTUs from upstream to downstream was defined as the action level for the St. Lawrence Sediment Removal Project during waterborne activities.

Real-time turbidity measurements were obtained from three monitoring locations, one 50 feet upstream of the western extent of the control system and two between 200 and 400 feet downstream of the eastern-most active installations, during the mobilization and installation of the Phase I sediment control system to evaluate any potential short-term effects of the operations. Measurements were collected near 50% water depth. Turbidity was also monitored if visible sediment releases were observed during sheet pile installations.

Reference

6.   "St. Lawrence River Sediment Removal Project Environmental Monitoring Plan." Prepared for General Motors Powertrain by BBL Environmental Services, Inc. May 1995.

4. Use of Acoustic Instruments for Estimating Total Suspended Solids Concentrations in Streams—The South Florida Experience (Patino et al., USGS, 2003)

An acoustic velocity meter (AVM) and an acoustic Doppler velocity meter (ADVM) were used in a study to estimate the total suspended solids (TSS) concentration in two southern Florida streams. The AVM system provides information on automatic gain control (AGC), which is an index of the strength of the acoustic signal recorded by the instrument as the acoustic pulse travels across a stream. The ADVM system provides information on acoustic backscatter strength (ABS), which is an index of the strength of return acoustic signals recorded by the instrument. Both the AGC and the ABS values increase as the concentration of suspended material increases.

The AVM system was installed in 1993 in the L-4 Canal, a man-made channel in northwestern Broward . The canal is approximately 40 feet wide and averages between 7 and 8 feet in depth. The water velocities in this canal range from –0.5 to 2.5 feet per second. The ADVM system was installed in 1997 in the North Fork Stream (a tidal channel), located in Veterans Park in southeastern Florida. The stream is about 280 feet wide and averages 8 feet in depth, with water velocities that range from about –1.5 to 1.5 feet per second and a salinity that varies from fresh to brackish (0.2 to 15 mg/l).

Depth integrated samples for TSS were collected at the L-4 Canal site using a DH-59 sampler and equal discharge increment (EDI) methodology, and samples at the North Fork site were collected using a point sampler at the same depth as the ADVM system and located 9 feet away from the transducer faces (near the start of the sampling volume). TSS concentrations ranged from 22 to 1,058 mg/l at the L-4 Canal site, and from 3 to 25 mg/l at the North Fork site.

Regression analysis techniques were used to develop empirical and site-specific relationships between the AGC and ABS results and the TSS and the two sites. The equation below describes those relationships:

$$TSS = 10^{\{A*[a + b* \log (salinity) + C * \log (temperature)] + d * \log (velocity) + e\}}$$

The relationships obtained using the site-specific equations produced good correlations, with coefficients of 0.91 and 0.87 at the L-4 Canal and North Fork sites, respectively. The results suggest that this technique is feasible for estimating TSS concentrations in streams using information from acoustic instruments.

Reference

Patino, E.; Byrne, M. J. "Use of Acoustic Instruments for Estimating Total Suspended Solids Concentrations in Streams—The South Florida Experience," U.S. Geological Survey, Ft. Myers, FL. Available at **http://water.usgs.gov/osw/techniques/TSS/Patino.pdf**, downloaded Feburary 2003.

5. Section K.6.2 – Correlation Analysis found in Appendix K: Water Quality Monitoring Pre-Design Field Test Dredge Technology Evaluation Report, New Bedford Harbor Superfund Site (USACE, 2001)

A Pre-Design Field Test was undertaken in order to evaluate the performance of a dredge system under consideration for use at the New Bedford Harbor Superfund Site. The objectives of the test are focused on the performance of the dredge system; this report segment evaluates the impacts on water quality associated with the test, including the following:

- Predictive modeling used to aid in the design of the water quality monitoring field program and to assess the utility of modeling for the full-scale remediation effort.

- Field monitoring to assess sediment resuspension during the dredging operation, to collect water samples for laboratory analysis, and to ground-truth the predictive modeling.

- Laboratory analysis of water samples (TSS, PCBs) to assess water quality impacts.

- Correlation assessment between the field and laboratory data.

Three correlation studies were performed on the data obtained from the monitoring samples:

- TSS vs. total particulate PCBs – Analysis of the data revealed an excellent correlation between the two parameters. The study yielded a coefficient of fit for the linear relationship of 0.84, suggesting that TSS serve as a good indicator of the particulate PCB concentrations associated with operations similar in scope to the pre-design work.

- Total particulate PCBs vs. total dissolved PCBs – Analysis of the data yielded a poor correlation between these parameters. An exponential function provided a better fit to the data.

- TSS vs. total dissolved PCBS – Analysis of the data provided a poor correlation between these parameters. An exponential function provided a better fit to the data.

A review of the individual dissolved/particulate data pairs indicated the following:

- For the reference samples, the dissolved and particulate PCB concentrations were generally similar on a per liter basis, with the dissolved sometimes exceeding the particulate.

- For the samples impacted by the dredging operations, the total particulate PCB concentration was generally increased to a much greater degree than the dissolved PCB concentration.

Analysis of the monitoring data also suggested the following:

- A moderate correlation between the total suspended solids measured in the lab and the turbidity measured in the field. The linear coefficient of fit for these data was 0.56. Measurement of both parameters from the same water parcel would be expected to increase the strength of the correlation.

- Given the different correlations indicated by the data, turbidity to TSS and TSS to PCB, the results suggest that field measurement of turbidity could be used as an indicator of the mobilization and transport of particulate-bound PCBs during the full-scale remediation activity.

Reference

USACE. 2001. "Appendix K: Water Quality Monitoring Pre-Design Field Test Dredge Technology Evaluation Report, New Bedford Harbor Superfund Site," *Pre-Design Field Test – Dredge Technology Evaluation Report*, New Bedford Harbor Superfund Site, New Bedford, Massachusetts. Prepared by Foster Wheeler Environmental Corporation, Boston, Massachusetts. August 2001.

6. Suspended Solids Flux Between Salt Marsh and Adjacent Bay: A Long-term Continuous Measurement (Suk et al., 1999)

The goal of this study was to establish an improved methodology to determine the suspended solids flux between Schooner Creek, NJ, a tidal salt marsh, and Great Bay, adjacent to it. The most significant difference in methods used in this study was related to data collection. Field data were collected continuously from March to October 1996.

A suite of instruments, including a current velocity sensor, a turbidity sensor, an automatic water sampler, a pressure transducer, and a data logger were placed in (and around) a location 300m from the mouth of Schooner Creek, to measure the velocity, water surface elevations, and suspended solids concentrations of the creek. Water velocity was measured at a depth corresponding to the mid-depth of the creek at high tide. The instruments were placed in the water on the deeper side of the creek so that they would remain submerged.

Total suspended solids (TSS) in the stream were quantified using turbidity as an indicator. A feasibility study performed prior to the experiment's initiation that examined 593 water samples over 25 different time periods found that that the measured suspended solids concentrations were statistically related to the measured turbidity. The average correlation coefficient for flood and ebb time periods averaged 0.827, indicating that turbidity measurements would provide surrogate measurements of the suspended solids concentration.

The water flux rate was derived from measurements taken by the submerged instruments, and calculated as a product of the current velocity and the area of the wetted cross section, and cumulative flow volumes were calculated using the average flow rate for successive time intervals.

The TSS flux was calculated as the product of the water flux and the TSS concentration. Two TSS fluxes were calculated:

- TSS fluxes for the entire recording period (periods of balance and imbalance) using TSS concentrations derived from the overall regression relationship.

- TSS fluxes for periods of time where the calculated water fluxes were more balanced, yielding net flux values that were not strongly impacted by a water imbalance.

Analysis indicated that the flow data are not continuous, and there are several different natural and artificial factors that may attribute to a water imbalance, though the researchers decided that net water import or export during a particular time was most likely due to the measurement of an incomplete cycle of water exchange across marsh boundaries other than the creek mouth.

The study also calculated a minimum number of water sample sets needed to produce a reasonably good TSS-turbidity regression relationship. To do so, varying combinations of water sample sets were used to develop a number of different regression relationships. The regression relationships were then used in the flux calculations, and the relative error was calculated.

The following observations were produced from the study:

- Data analysis indicated that the cumulative and cycle fluxes calculated for the entire recording period are considerably uncertain due to an imbalance in the calculated water fluxes.

- Data analysis indicated that the coefficient of correlation between the cumulative TSS fluxes per tidal cycle and the average TSS concentration differences was 0.71. The flow-weighted average TSS concentration resulting from all of the water balance periods during the flood tide was higher than that during the ebb tide, contributing to a net import of TSS.

- Data suggested that, for this study, a reasonably good overall TSS-turbidity regression was established when five data sets with correlation coefficients greater than or equal to 0.80 were used.

Reference

Suk, N. S.; Guo, Q.; Psuty, N. P. "Suspended Solids Flux Between Salt Marsh and Adjacent Bay: A Long-term Continuous Measurement," *Estuarine, Coastal, and Shelf Science,* Vol. 49, pp. 61-81, 1999.

Attachment F-2

PCB Analytical Methods
Detection (Reporting) Limits in Water

Attachment F-2
PCB Analytical Methods
Detection (Reporting) Limits in Water

1. **CLP Method OLM04.1** (September 1998)
   Contract-required quantitation limit is 1 μg/L for all Aroclors
   (CRQL for Aroclor 1221 is 2 μg/L)
   Laboratories can report lower detections (e.g., 0.5 J [μg/L])

2. **SW-846 Method 8082** (Rev 0, December 1996)
   MDLs (method detection limits) for Aroclors range from 0.054 to 0.90 μg/L
   (Method provides no data as to Aroclor-specific MDLs)

3. **PCB Congeners - Dual Column GC/ECD** (Laboratory-specific)
   STL/Colchester Vt (formerly Aquatec)
   Detects inidividual PCB congeners at a detection limit of 0.001 μg/L
   (Monochlorobiphenyls at 0.005 μg/L)
   (Other labs have other methods with varying detection limits)
   [Need to verify current STL limits - post-Hudson RI/FS - as used for Reynolds, e.g.]

4. **NYSDEC Analytical Services Protocol Low-Concentration Method (91-6)**
   CRQL is 0.2 μg/L for Aroclors except for 1221 (0.4 μg/L)
   (There is probably a corresponding USEPA CLP method, which I don't have)

5. **USEPA Method 505**, Revision 2.1 - 1995 (Organohalide Pesticides and PCBs by microextraction/GC)
   MDL for Aroclors 1016, 1248, 1254 - about 0.1 μg/L
   MDL for Aroclor 1260 - about 0.2 μg/L
   MDL for Aroclor 1242 - about 0.3 μg/L
   MDL for Aroclor 1232 - about 0.5 μg/L
   MDL for Aroclor 1221 - about 15.0 μg/L
   (from Method 505 Revision 2.0, USEPA EMSL, 1989)

6. **USEPA Method 508**, Revision 3.1 (1995). Determination of Chlorinated Pesticides in Water by GC/ECD.

- Note to method summary states that the extraction is similar to Method 608 (q.v.), and the extract can be analyzed by 508, 525, or 608; however, no performance data for Aroclors were collected as part of method development for 508.

- EDLs (reporting limits) for most single-component pesticides are in the 0.01 μg/L to 0.05 μg/L range (a few are higher and a few are lower).

- This method is supposedly being used by Waterford for monitoring its drinking water supply. The detection and reporting limits would have to be developed on a laboratory-specific basis. Muliti-component analytes (such as Aroclors, and also toxaphene and chlordane) typically have higher reporting limits than single-component pesticides.

7. **USEPA Method 680** (PCBs by GC/MS)
   Arocolor detection limits are about 100 μg/L

8. **USEPA Method 608** (Pesticides/PCBs by dual column GC)
   Aroclor Detection limits 0.5 μg/L (1.0 μg/L for Aroclor 1221)

9. **USEPA Method 525.2** (1995 revision)
   Method uses solid/liquid extraction by either disk or cartridge; and analysis using quadropole MS or ion trap. MDLs are presented for method analytes for each of the four possible combinations; except Aroclor MDLs only by disk and ion trap. Sensitivity is better for more chlorinated arolcors. MDLs range from 0.018 μg/L for 1260 to 0.054 μg/L for Aroclor 1221.

10. **USEPA Method 1668A** (December 1999) - Chlorinated Biphenyl Congeners in Water, Soil, Sediment, and Tissue by HRGC/HRMS.
    - Detection limits (EMDLs) and reporting limits (EMLs) are provided for more than 150 congeners in both water and non-aqueous matrices.
    - Method is more sensitive for less-chlorinated congeners.
    - Reporting limits for individual congeners range from 50 to 1000 pg/L (10 pg/L for BZ#2) in water (detection limits [EMDLs] are typically 1/3 to ½ the reporting limit [EML]).
    - Reporting limits range from 5 to 100 ng/kg (except 1 ng/kg for BZ#2) in non-aqueous samples (detection limits [EMDLs] are typically 1/5 to ½ the reporting limit [EML]).

11. **Green Bay Method.** Original method not reviewed (or obtained). *Not included in the GE August Design Support Sediment Sampling and Analysis Plan (Revision 1, August 2002).* Reportedly a single-column PCB congener GC/ECD method.

General notes on units of measure:
- g/L = parts per thousand (10^{-3});
- mg/L = parts per million (10^{-6});
- μg/L = parts per billion (10^{-9});
- ng/L = parts per trillion (10^{-12});
- pg/L = parts per quadrillion.(10^{-15}).

Attachment F-3

**Memo Regarding PCB Analyses; Whole Water Extracts vs. Separated
Particle and Filtrate Extracts**

February 25, 2003

To: Kelly Robinson, Earthtech (TAMS)
From: Richard Bopp, RPI
Re: PCB Analyses; Whole Water Extracts vs Separated Particle and Filtrate Extracts

Background

Since I first analyzed Hudson River water samples for PCBs in the late 1970s, I have
been interested in particle/water partitioning. Consequently, I have always filtered the
samples and extracted and analyzed the particles and filtrate separately. In addition,
based on considerations of analytical sensitivity, I have always analyzed large volume
(typically 18 liter) water samples. These procedures were adopted by the USEPA for the
water column PCB samples that we collected and processed as part of the Hudson River
PCBs Reassessment.

Several other important datasets rely on an EPA-approved whole water extraction and
analysis of much smaller volume (typically 1 liter) samples. These include

- The USGS monitoring in the upper Hudson. This program provides the longest
  historical record of water column PCB levels.

- The GE monitoring between Rogers Island and Schuylerville conducted under
  consent order with the NYSDEC as part of the remnant deposits monitoring
  program. This set of samples, collected approximately weekly since 1997
  provides, by far, the most detailed picture of PCB transport ever developed (J.
  Tatten, Master's Project, RPI, 2000; Task 3 Final Report to NYSDEC, Contract
  C003844, 2000).

In 1993 I was at RPI and supervising the collection and processing of the water column
samples for the Hudson River PCBs Reassessment. As I recall, I suggested that on one of
the transects we collect duplicate samples for PCB analysis through NYDSEC at the
NYSDOH labs. In addition, since their standard procedure was whole water extraction, it
was arranged that at least some of the samples also be analyzed as separate particle and
filtrate fractions. This would allow a more direct comparison with the EPA sample
analysis and provide a test of my general impression that whole water extraction would
not be particularly efficient at recovering particle-associated PCBs. The suggestion was
welcomed at NYSDEC and collaboration was facilitated by the fact that I had been
employed there in 1990-91.

Analysis and interpretation of the data from this exercise was to form the basis of the
Master's project of Christine Juliano. After an initial data gathering and analysis effort,
Christine decided to work on a different project and completed her Master's. My
preliminary look at the data indicated that whole water extraction missed a significant
fraction of the particle-associated PCBs. Although based on very limited data, I have

used this observation often to support my geochemical bias toward separate particle and filtrate extraction and analysis.

Over the past month, I have had two requests for a more quantitative assessment of this data. Both were related to water column monitoring associated with the proposed dredging. The first was from Kelly Robinson at Earth Tech (TAMS), the primary EPA contractor on the Upper Hudson River PCB project. A few days later, Roger Sokol of the NYSDOH requested similar information specifically for monitoring the Waterford, NY drinking water supply and raw water intake on the Hudson. I was able to locate files prepared by Christine Juliano that contained water column PCB data from the upper Hudson consistent with events described above.

More Detailed Information

The sample ID format and numbering used in the files indicates that the samples were collected during EPA transect 4 (April 12 to April 14, 1993) at Stillwater (0007), Waterford (0008), the Hoosic River (0012), Mohawk River (0013), and Green Island Bridge (site 0014). Two of the samples, Waterford and Green Island, have data for whole water and separate particle and filtered water analyses. Further confirmation of the identification of these samples comes from the fact that the TSS levels in the files prepared by Christine Juliano are identical to those reported for samples TW-0004-0008 (34.0 mg/l) and TW-0004-0014 (39.8 mg/l) in the EPA Database. More specific collection information can most likely be retrieved from the detailed field notes kept by Rensselaer personnel and submitted to TAMS a part of the official record of our work with EPA on the reassessment. The rest of this report will refer to the Waterford (004-0008, 04/13/93) and Green Island (004-0014, 04/13/93) samples.

As I recall, I was informed that the separation of particulate and dissolved phases for the NYSDOH analysis was accomplished by pouring the water sample through a soxhlet extraction thimble. This simple procedure should be comparable to separation by more standard filtration techniques that typically employ pre-fired glass fiber filters. The corresponding EPA samples that we collected were filtered by Kevin Reed of RPI through pre-fired Whatman GF/F filters. Soxhlet extraction thimbles used in PCB analyses are also treated to minimize blanks. Paper thimbles are typically pre-extracted and glass fiber thimbles are pre-fired.

Results

- In terms of total PCBs, the DOH values reported for the whole water extracts were about half of the (particulate + dissolved) PCBs in the replicate samples (Table 1).
- At the congener level, whole water extraction yielded results lower than (P + D) in every case with only one exception (BZ 24, 27). Figures 1 (Waterford) and 2

(Green Island) present data for a range of more abundant congeners that together comprise over half the total PCBs.

- The figures also show that the differences between whole water and (P + D) results tend to be less for the lower chlorinated congeners. This is consistent with a simple model of the whole water extraction process – complete recovery of dissolved PCBs and less efficient recovery of particulate phase PCBs.
- Based on this first order model applied at the congener level, the whole water extraction missed $61 \pm 20\%$ of the particle-associated PCBs in the Waterford sample (Table 2) and $72 \pm 13\%$ in the Green Island sample (Table 3).

Implications

- The above analysis provides support for the logical assumption that whole water extraction will result in an underestimate of total PCBs. It is also logical to assume that the degree of under-recovery would depend significantly on the details of the procedure (the number of extraction cycles, the solvent used, the percentage of solvent removed between extraction cycles, the degree of sample agitation etc.).
- If the simple model presented above is applied, the degree of under-recovery will also depend on the TSS in the sample. Using an average particle extraction efficiency of 33% (based on the DOH analyses) and an average upper Hudson PCB particle/water distribution coefficient of 10^5 (Bopp et al., Final Report to NYSDEC, Contract C00708, 1985), first-order error estimates can be made.

TSS (mg/l)	% of PCB on Particles	% under-recovery of total PCBs
2	17	11
10	50	33
40	80	53
100	91	61

- This analysis raises the possibility that historical (USGS) estimates of PCB transport in the upper Hudson that focused on high flow, high TSS, high transport events may be low by on the order of 50% and suggests a low bias to any transport estimates that utilize the weekly GE water column monitoring data.
- The potential for significant under-recovery of PCBs when using whole water extractions should be considered in the design of any future monitoring program.

Cc: Roger Sokol, NYSDOH

Tables

Table 1. 'Total PCBs in samples collected April 12 -14, 1993 (all PCB concentrations in ng/l)

	Waterford DOH	(0008) EPA	Green I DOH	(0014) EPA
Particulate	225.4	159.8	227.7	144.5
Dissolved	74.4	75.0	50.9	53.5
P + D	299.8	234.8	278.6	198.0
Whole Water	159.9		110.6	
TSS (mg/l)	34.0		39.8	

Table 2. 'Waterford

CONGENER	004-0008 Whole Water	004-0008 Particulate	004-0008 Filtered Water	sum P+F		%P missed		%T missed
BZ-10,BZ-4	17	9.5	13	22.5		58		24
BZ-19	4.8	2.3	3.6	5.9		48		19
BZ-18	12	5.1	6.9	12		0		0
BZ-15,BZ-17	11	8.5	5.8	14.3		39		23
BZ-16,BZ-32	4.3	3.5	2.8	6.3		57		32
BZ-31	7.5	12	4.2	16.2		73		54
BZ-28	8.3	14	4.6	18.6		74		55
BZ-20,BZ-33,BZ-53	3.6	5.6	2.1	7.7		73		53
BZ-52	6.4	9.5	3	12.5		64		49
BZ-49	5.6	8.8	2.2	11		61		49
BZ-47	4.4	7.6	1	8.6		55		49
BZ-44	3.7	6.2	1.5	7.7		65		52
BZ-37,BZ-42,BZ-59	1.2	3.2	1	4.2		94		71
BZ-41	3.1	5.3	1.2	6.5		64		52
BZ-70	4.8	11	1.6	12.6		71		62
BZ-66,BZ-95	7.8	18	2.1	20.1		68		61
BZ-110,BZ-77,BZ-136	2.3	6.4	*0.5*	6.9		72		67
Totals	107.8	136.5	57.1	193.6		61		45
					Std. Dev.	20	Std. Dev.	19
BZ-24,BZ-27	7.7	2.7	4	6.7		-37		-15

Table 3. Green Island

CONGENER	004-0014 Whole Water ng/L	004-0014 Particulate ng/L	004-0014 Filtered Water ng/L	sum P + F	%P missed	%T missed
BZ-10,BZ-4	13	8.5	9.9	18.4	64	29
BZ-19	3.8	1.7	3	4.7	53	19
BZ-18	7.9	5.1	5.2	10.3	47	23
BZ-15,BZ-17	7.4	8.9	4.1	13	63	43
BZ-16,BZ-32	3.3	3.5	2.2	5.7	69	42
BZ-31	5.3	13	2.5	15.5	78	66
BZ-28	5.9	15	2.7	17.7	79	67
BZ-20,BZ-33,BZ-53	3	5.7	1.7	7.4	77	59
BZ-52	4.6	8.6	2.1	10.7	71	57
BZ-49	4.1	8.6	1.5	10.1	70	59
BZ-47	3.2	6.7	*0.5*	7.2	60	56
BZ-44	3.7	6.2	1.5	7.7	65	52
BZ-37,BZ-42,BZ-59	1	3.1	1	4.1	100	76
BZ-41	2.2	9.4	1	10.4	87	79
BZ-70	3.3	10	1	11	77	70
BZ-66,BZ-95	5.3	15	1.1	16.1	72	67
BZ-110,BZ-77,BZ-136	1.7	6.4	1	7.4	89	77
TOTALS	78.7	135.4	42	177.4	72	55
					Std. Dev. 13	Std. Dev. 18
BZ-24,BZ-27	5.2	2.3	2.8	5.1	-4	-2

Figures

Figure 1
WATERFORD
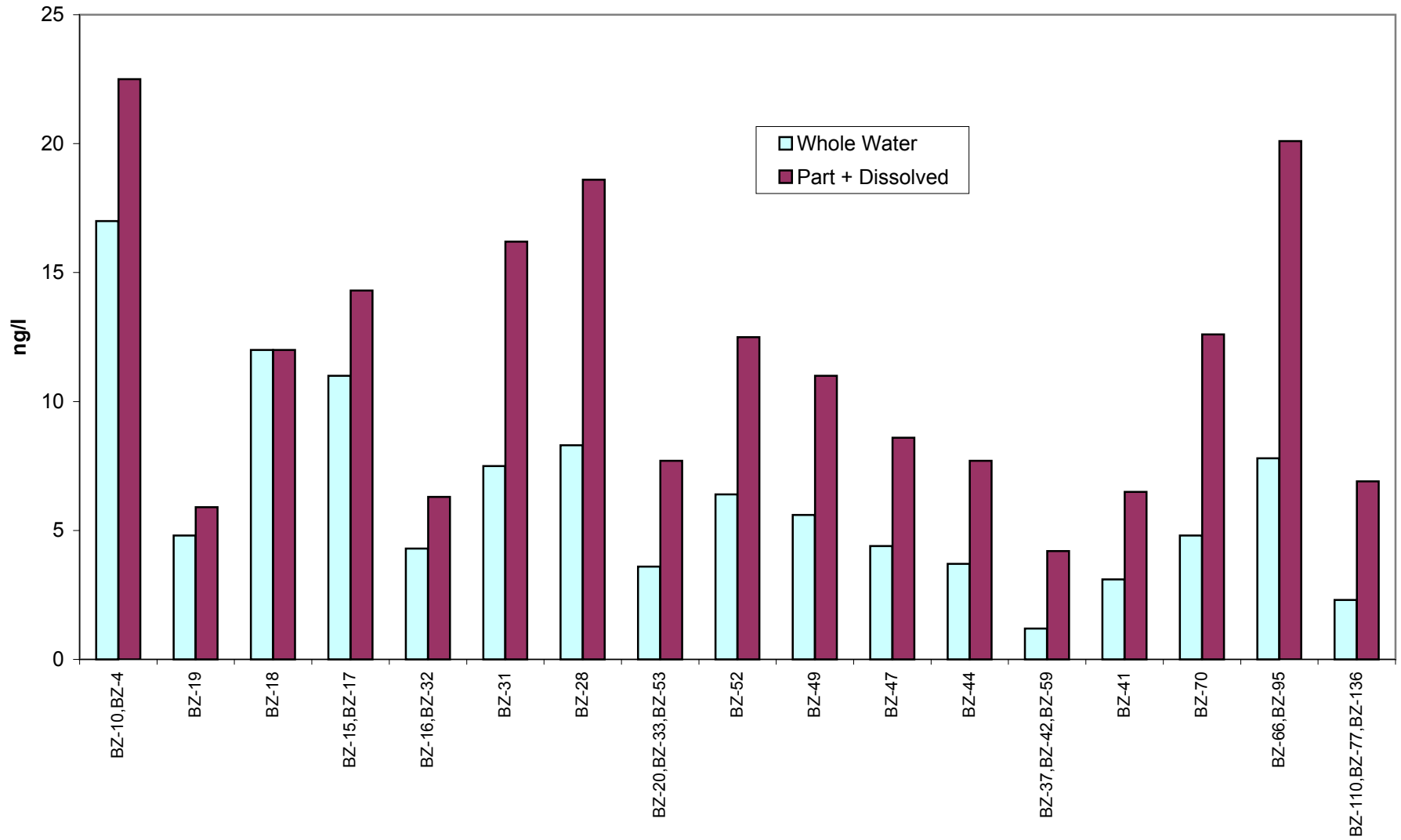
Figure 2
GREEN ISLAND
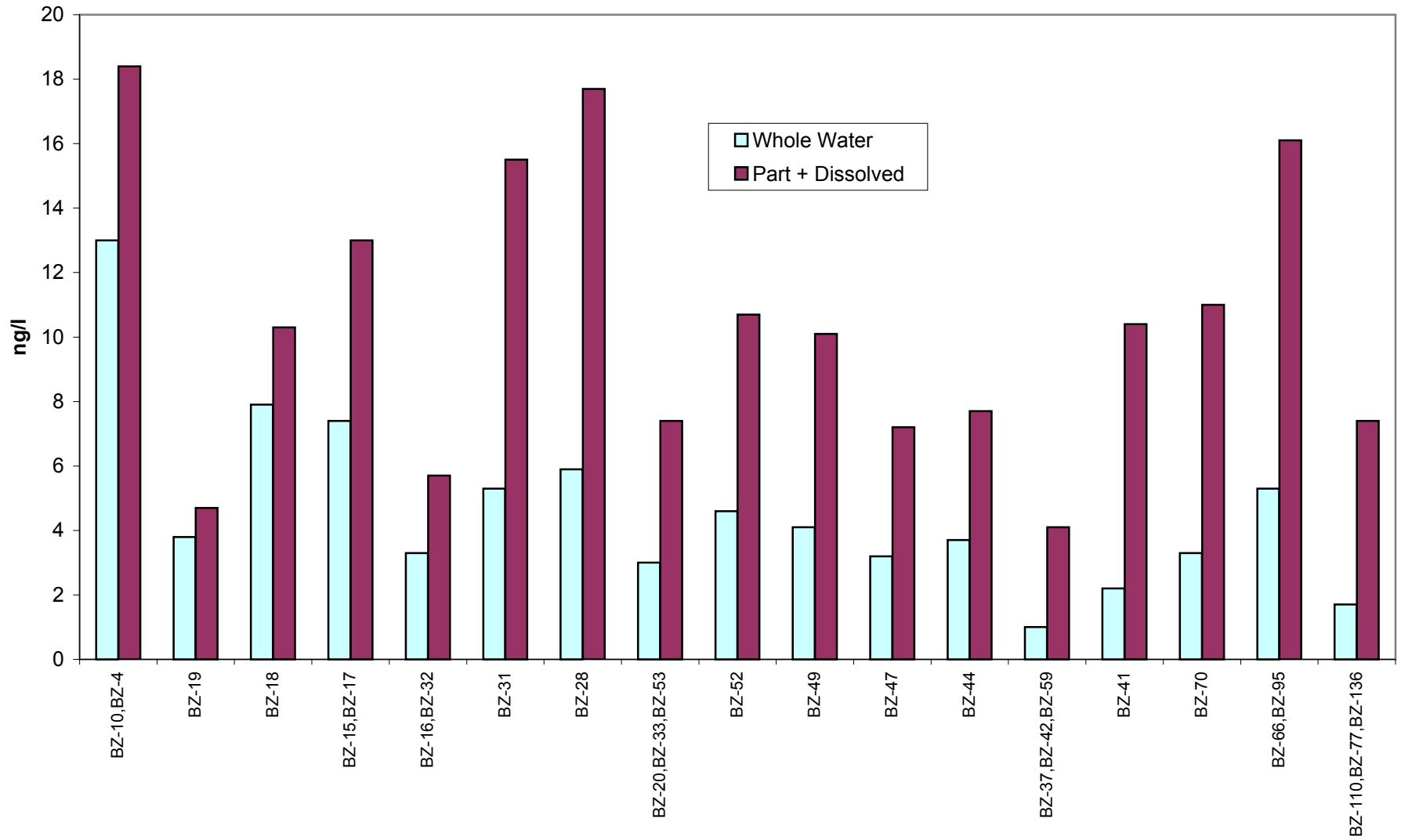
Attachment G

A Discussion of Data Quality Objectives

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A Discussion of Data Quality Objectives

Introduction

The monitoring plan for the Resuspension Performance Standard is summarized in Tables 1-2, 1-3 and 1-4 of the main document. The main objectives of the monitoring plan are described in Sections 1 to 3 below along with the techniques intended to satisfy these objectives. Additional discussion on rationale for the sampling program can be found in Section 2 of the standard. This analysis represents an initial analysis of the data quality objectives that will undergo subsequent refinement in the preparation of the quality assurance plans for dredging-related monitoring. As such, it is expected that the monitoring requirements developed for the standard represent a minimum level of monitoring and that additional sampling beyond these requirements will be needed to completely understand the nature of any dredging related release. These requirements are primarily intended to document compliance with the various criteria of the Resuspension Performance Standard while also providing some initial information on the mechanisms of PCB release. In addition to determining compliance with the resuspension criteria, the monitoring plan includes elements that serve to verify assumptions made about the behavior of the contaminant releases due to dredging (*e.g.,* PCB dissolution, suspended solids settling and dissipation). Information collected to verify these assumptions during the Phase 1 period should serve to improve the monitoring program during Phase 2 in several ways. The Phase 1 data should permit the identification of the most effective monitoring locations and monitoring techniques as well as identify any that are not useful. This information should also permit a reduction in the frequency and complexity of monitoring during Phase 2.

In Sections 1 to 3 of this attachment, the main data quality objectives are identified followed by a discussion of the sampling approaches needed to satisfy each objective

Section 4 of this attachment describes the adequacy of the sampling frequencies required as part of the routine and non-routine monitoring programs. These are derived using USEPA defined methods for assessing statistical uncertainty (USEPA, 2000). The analyses cover only routine monitoring and the minimum levels of contingency monitoring as defined in the Resuspension Standard. Additional monitoring related to the required engineering studies at the Concern and Control Levels (as well as exceedance of the standard threshold) may be required, depending on the anticipated cause of the exceedance. The design of these additional monitoring programs may be developed during the remedial design period. Alternatively, *ad hoc* monitoring plans may be developed by the design team during the actual dredging operation in response to observations made at the time.

A particular limitation to the analysis presented in Section 4 of this attavchment is information on the variance of river conditions in response to dredging-related releases. Little data exist on which to develop the estimate of variance. As a result, the variation of

baseline conditions was used as a means to estimate the variance for dredging operations. These estimates for sampling requirements and the associated error rates will require review once additional data becomes available during Phase 1.

1   Objectives for Far-Field Monitoring in the Upper Hudson

Several data quality objectives are addressed by the far-field monitoring program in the Upper Hudson. This program is the primary monitoring effort for the protection of public water supplies and for determining the magnitude of long-term PCB releases. Following the statement of each data quality objective is a discussion of the sampling techniques to be used to satisfy the objective.

Objective I: Provide a set of data to demonstrate compliance with the PCB concentration components of the Resuspension Standard (*i.e.,* the 350 ng/L criteria for the Concern and Control Levels and the 500 ng/L criterion for the standard threshold).

- Dredging-related operations are expected to occur throughout the Upper Hudson between Ft Edward and Waterford. Hence dredging-related PCB release may occur over the entire region as well. In particular, while the majority of dredging is focused north of Schuylerville, boat traffic and other operations are expected to occur downstream of Schuylerville. Thus PCB concentrations must be monitored throughout the Upper Hudson. Additionally, PCB release due to dredging is not expected to be constant with time but is expected to vary substantively over time. Thus discrete grab samples collected at one station at one point in the day may miss more substantial release events occurring at other times. As the river carries these releases, natural mixing and dispersion will serve to homogenize PCB concentrations to some degree, spreading them out and making them easier to sample at locations farther downstream. Thus multiple stations provide a basis to capture conditions representing a longer period of time. Note that the desire to obtain many samples from the river to characterize conditions must be tempered by the availability of laboratories to analyze the samples. For this reason sampling under routine conditions (expected to be the majority of the conditions while dredging) will only require daily samples from the far-field stations plus a limited number of longer-term integrated samples (see Table 1-2 in Section 1 of the Resuspension Performance Standard). This consideration also recognizes the need to obtain and analyze samples sufficiently rapidly to address Objective II below. An alternative to these discrete samples is the collection of daily composite samples, integrated over a 24-hour period at each station. These samples still require the collection of a cross section composite for each day. Additional sampling may be required if 24-hour composites are collected when various action levels are exceeded and two phase sampling is required.

- It is necessary to correctly characterize the PCB concentration throughout the river's cross-section, recognizing that both baseline and dredging-related releases create heterogeneous PCB concentrations. This has been extensively demonstrated by the

paired sample data collected at TID-West and TID-PRW. For this reason, at least five points are required at each sampling station, based on equal area or equal discharge considerations as given by USGS guidance. Multiple points are required for discrete samples as well as the alternative daily composite samples.

- To support the use of discrete samples as representative of mean river conditions, it is also necessary to obtain integrated samples. These samples will serve to demonstrate compliance with the standard during periods between discrete samples. Integrated samples will cover one to two week intervals, providing a longer perspective on PCB transport and concentration with relatively little increase in the total number of PCB analyses. Rapid turnaround of results cannot be provided for the integrated samples since they take longer to collect; rather they provide confirmation of data obtained from daily discrete samples. As such they are needed during Phase 1 to provide supporting data for the discrete samples. If they confirm the viability of the discrete sampling program, these samples may be dropped or greatly reduced during Phase 2.

- Samples must be collected at sufficient frequency to provide a reasonable statistical certainty that conditions are in compliance with the Resuspension Standard criteria. Higher statistical uncertainty is acceptable when concentrations are well below the standard criteria. As the various action levels and the standard threshold are approached, sampling frequency must be increased to provide greater certainty that conditions are still in compliance. In particular it is important to minimize the false negative error, the error of accepting conditions to be in compliance when in reality they are not. The issue of sampling frequency is extensively discussed in Section 4 of this attachment.

Objective II: Provide a means to rapidly assess water column PCB levels so that the USEPA can advise public water suppliers when water column concentrations are expected to approach or exceed the federal MCL (*i.e.,* 500 ng/L) during the remediation. In this manner, public water suppliers can take contingency actions, if needed, to maintain safe water for their users. Appurtenant to this objective, determine the relationship of dredging-related PCB contamination at the upstream far field stations (TI Dam and Schuylerville) to that at the downstream far-field stations (Stillwater and Waterford) in order to use the far-field stations near the remediation as predictors of downstream concentrations entering the public water intakes.

There are several aspects of the monitoring plan that are required to achieve these closely related objectives. These are described below.

a) Measurements of PCB concentrations at all Upper Hudson far-field stations are needed on a daily basis to identify possible exceedances of the standard threshold and any action level criteria. These data satisfy both components of this objective, since the data will document the PCB concentrations and also serve as a database to resolve

the relationship between upstream and downstream PCB concentration increases related to dredging.

b) Reduced turnaround time for PCB samples from the two far-field stations nearest to the dredging operations is required. During Phase 1 these stations will probably be TI Dam and Schuylerville, although the Phase 1 dredging area has not yet been defined. The results from these stations will be used to assess the need to warn the public water supplies that the concentrations entering the intakes may be elevated. The travel time between remediation activities in River Sections 1 and 2 and the Waterford public water supply intakes is generally two days. Thus, in order to have information from the primary dredging areas in time to provide a warning to the downstream intakes, a turnaround time of 24 hours or less is required for the samples obtained from the two nearest downstream far-field stations.

c) While actual PCB measurements provide the most certain basis for assessing PCB loads and concentrations, these cannot be obtained in real time. A reliable real time indication of PCB levels can be derived from continuous or high frequency suspended solids monitoring. Resuspension of contaminated sediment is thought to be the primary mechanism of dredging-related contamination release. When verified, suspended solids monitoring provides one of the best means of warning the public water supplies of potential exceedances, since it provides the longest lead time between knowledge of the release and its arrival at the downstream intakes. Additionally, as the dredging operations move farther downstream, suspended solids monitoring will provide the only real time data for the protection of downstream impacts. Specifically in River Section 3, there will be insufficient time to collect, analyze and evaluate a PCB sample and still warn the downstream intakes. As a result, frequent sampling of suspended solids and other indicator measures will be required in the standard to develop this capacity.

Objective III: Provide a set of data to demonstrate compliance with the PCB load components of the Resuspension Standard (*i.e.,* 300 g/day and 600 g/day).

a) Just as for Objective I, dredging-related operations are expected to occur throughout the Upper Hudson between Ft Edward and Waterford, increasing PCB loads as well as concentrations. PCB loads, however, represent a longer-term concern since their impacts will take longer to occur and require a sustained level of loading in order to have an impact. Nonetheless, it is desirable during Phase 1 to identify substantive increases in load soon after they occur so that the root cause can be identified. To this end, the monitoring frequency required to satisfy the concentration criteria will also satisfy this objective.

b) Since PCB loads over time are the primary concern of this objective, it is desirable to obtain integrative samples for this objective as well. For this reason, integrative samples will be obtained at the four main far-field stations during Phase 1 as

discussed under Objective I. These will provide confirmation of the initial conclusions drawn regarding PCB loads based on the more frequent discrete samples.

c) Data on river discharge is also needed to address load considerations. Data from the USGS stations at Ft Edward and Waterford will be used to this end. In the event that the USGS discontinues these stations, data on flow must be obtained by an alternate means. Additional data on meteorological conditions must be obtained to supplement the USGS data and permit an accurate representation of flows at the stations not monitored by the USGS.

d) Sample collection must be timed to capture the impacted water column. If samples were collected each day from the nearest far-field station at the onset of the operations, it is unlikely that the water collected would show the dredging-related impacts. The plume will widen and lengthen as it travels downstream, making it more likely that the downstream stations will capture dredging-related impacts. (This is not, however, a time-of-travel sampling. Although the parcel of water sampled must be impacted, but the same parcel of water need not be tracked as it passes down the river.)

e) Equal discharge increment (EDI) or equal width increment (EWI) sampling as defined by USGS will be required. This type of sampling method is required to capture a representative cross-sectional sample. A single center channel station will not be sufficient, because there may not have been sufficient mixing across the channel and plumes confined to the shoreline by river hydrodynamics would not be measured, resulting in low-biased results.

Objective IV: Determine the primary means of PCB release via dredging-related activities. (Verify that dissolved phase releases are minimal as estimated by modeling and that the primary mechanism of release is suspension of sediment.)

a) During the public comment period on the Hudson River ROD, concerns were raised that dredging of PCB contaminated sediment could release a substantial amount of dissolved phase PCBs. Calculations to determine if and how such a release could occur (Attachments C and D) have indicated that this scenario cannot occur and that the primary release mechanism would be resuspension of contaminated sediment. This mechanism would be accompanied by an increase in suspended solids concentration and could be tracked in the near field.

b) Though convincing, the calculations done to determine the primary mechanism of release need to be verified in order to be certain that the goals of the ROD can be achieved (long-term recovery of the river, protection of the environment and human heath). This will be accomplished by split phase sampling for PCBs at the two far-field stations closest to the dredging operations. PCB concentrations at these locations are less likely to be at equilibrium between dissolved and suspended fractions and can provide information regarding the nature of the released PCBs. Beyond these

locations, equilibration between phase is likely to exist, thereby reducing the value of split phase analysis. At these stations, whole water PCB analysis will continue.

c) Additional parameters will be required to aid in the interpretation of these split phase data. Dissolved organic carbon, suspended organic carbon, suspended solids and temperature provide an indication of the distribution of dissolved phase and suspended phase PCBs. These parameters will be measured for routine monitoring and contingency monitoring. In this manner, changes in these supplemental parameters may help identify the nature of the release mechanism responsible for the PCB release.

d) Resolution of this concern is only necessary in the event that PCB loads and concentrations exceed the action levels or the threshold criterion. Thus split phase sampling will only be required in response to these exceedances. Split phase sampling is not required under routine sampling.

Objective V: Determine the baseline PCB levels entering River Section 1 from upstream sources.

a) PCBs entering River Section 1 should be identified to differentiate these additional concentrations from the release from baseline and dredging-related contributions. Based on monitoring data from the past five years, PCBs have been not detectable or at low concentrations entering River Section 1. However, changes in upstream conditions (such as construction at the source areas) could result in higher PCB concentrations entering the TI Pool. Low frequency monitoring at Bakers Falls and Rogers Island for PCBs will be required for this purpose. If the contribution from upstream sources were to increase, the Bakers Falls and Rogers Island results should document this and will be used to adjust the dredging-related load contribution. This information would help to avoid an unnecessary enforcement of the engineering or monitoring contingencies of the standard and would be done on a case by case basis. With USEPA's approval, the frequency at Rogers Island may be further reduced if these concentrations are shown to be consistently low relative to dredging-related releases.

b) Both Bakers Falls and Rogers Island stations are needed for this purpose. An important assumption in the ROD was the continued reduction of the releases from the GE Hudson Falls facility. Differences in PCB concentration and load between these two stations will be used to document this process. In the event that these data are collected as part of other remedial activities upstream of Rogers Island, these data do not have to duplicated by the dredging-related monitoring. However, these data must meet the data quality objectives defined here and in the subsequent quality assurance plans issued for the Resuspension Performance Standard.

c) Detection limits for Total PCBs for these data must achieve the same detection limits as used for the far-field monitoring program, approximately 0.5 ng/L.

d) Additional data will be required to aid in the interpretation of downstream data. Baseline levels of dissolved organic carbon, suspended organic carbon, suspended solids and temperature are needed to characterize the changes in these parameters that may be caused by dredging-related activities.

e) Since baseline conditions should not change in response to dredging-related releases, the frequency of baseline monitoring does not increase in response to action level or threshold exceedances.

Objective VI: Determine ancillary remediation-related effects on the river (e.g., barge traffic related resuspension, spillage during transit or off-loading of sediment) that may occur in areas that are not captured by the nearest representative far-field station.

1. During Phase 1, the remediation will probably be limited to the TI Pool. Once the material is dredged it will be conveyed to another location for further processing and shipping to a landfill. This destination may not be in the TI Pool, resulting in transport of contaminated material throughout stretches of the Hudson River by barge or pipeline. To verify that the transport of material is not causing the release of PCB contamination to an extent that would cause exceedance of the resuspension criteria, sampling will be required at each Upper Hudson River far-field station (except Bakers Falls) at least once per day.

Objective VII: Verify that the water column PCB concentrations developed from the grab samples adequately characterize the average concentration.

1. Discrete grab samples will be used for comparison to the PCB flux and concentration resuspension criteria. The Resuspension Performance Standard requires that samples must be timed to capture the impacted water column, increasing the likelihood that the samples will be representative of the dredging-related impacts. As described in Section 4 of this attachment, the sampling frequency is sufficient to compare the results of the analyses to the resuspension criteria with confidence, but this analysis is based on an assumption of the variability of the water column concentrations. This estimate of variability is derived from the baseline conditions, which do not include the added variability of the dredging-related releases. This added variability could change within a day as different operations are completed and different dredge operators are employed. To verify that the grab samples are sufficiently indicative of average river conditions, integrating samplers are required for deployment periods ranging from two weeks under routine monitoring to one day under Control Level monitoring.

2. Integrating samplers are also required for daily measurements in place of discrete grab sampling at Stillwater and Waterford at the Concern and Control Level monitoring as well. This sampling method is used because of the concern that the water column concentrations are approaching the MCL. Integrating samples are used

here instead of multiple grab samples to reduce the overall number of PCB analyses while still obtaining data on PCB concentrations over a 24-hour period.

Objective VIII: Confirm the exceedance of the various action level criteria as well as the standard criterion.

1. Sampling frequency must be increased to verify exceedances of the resuspension criteria. At lower levels of exceedance, the consequences of error in deciding if the resuspension criteria have been exceeded are less serious than at higher levels of exceedance. Hence a higher level of decision uncertainty is acceptable at exceedances involving the lower action levels. At the Evaluation Level, the concern is adherence to best practices and long term PCB release impacts, concerns that do not require a rapid (*i.e.,* 24 hour) response. At PCB concentrations close to or above the Resuspension Standard, public water supplies could be impacted and a shutdown of the dredging operations may be required. Thus, a greater level of certainty is required when the consequences are greater. This is a primary reason for requiring increased frequency of sampling in the standard. The development and level of certainty provided by the various sampling regimes are further discussed in Section 4 of this attachment.

2. An increase in monitoring frequency will be required as a contingency at the two representative far-field stations during Phase 1. These stations provide the best opportunity to document river conditions in response to dredging-related releases and also provide a warning to downstream public water supply intakes. With the uncertainty related to dredging-related releases, the second station will confirm the observations of the nearest far-field station and thus provide as sound basis for whatever response actions are required.

3. Monitoring of the downstream far-field stations (Stillwater and Waterford) for PCBs will be changed to daily integrated sampling to capture the average concentrations that would be entering the public water supply, while PCB concentrations collected from stations nearer to the remediation may be approaching the MCL. Data from the integrated far-field samples provide further subsequent confirmation of the estimated concentrations based on conditions closer to the dredging operations. Results from these downstream stations can be used to refine the means of predicting the PCB concentrations that will enter the public water supplies based on the concentrations measured nearer to the remediation. These results will indicate the degree to which the PCB concentrations dissipate as the water column passes downstream. The switch from a daily discrete sample to an integrated sample reflects the need to characterize the entire day's water conditions while minimizing the number of samples collected, so that results can be rapidly available and interpreted.

2   Objectives for Monitoring in the Lower Hudson

Objective IX: Determine the extent of short-term impacts to the Lower Hudson River and examine the effect of Upper Hudson dredging activities on Lower Hudson PCB concentrations.

1. The monitoring program for the Lower Hudson is designed to measure the short-term impacts to the freshwater portion of the river (previously referred to as the Mid-Hudson River during the Reassessment) resulting from the remediation. The sampling requirements in the Lower Hudson are not designed for comparison to the resuspension criteria. This is addressed by the frequent sampling at Waterford, which will be extrapolated to conditions downstream. Requirements for additional monitoring at the public water supply intakes will be prepared as part of the community health and safety plan (CHASP) currently under development. The Lower Hudson stations are intended to characterize general water column conditions in response to elevated PCB concentrations and loads originating from dredging. These stations consist of a single center channel location that can be readily reoccupied. Cross sectional sampling is not required, since flow is not unidirectional and thus flux cannot easily be estimated.

2. The frequency of sampling is increased in the Lower Hudson in response to greater loads and concentrations in the Upper Hudson, specifically, when Troy is expected to exceed 350 ng/L. This is done to examine Lower Hudson conditions in response to these loads as part of the documentation of the recovery of the river.

3   Objectives for Suspended Solids Monitoring in the Upper Hudson

Objective X: Provide a real time indication of suspended solids release in the near field.

1. A real time indication of the amount of suspended solids in the water column in the near field will aid the dredge operators in minimizing the release of suspended solids and associated PCBs during the remediation. These monitoring will also provide the earliest evidence for a substantive PCB release and allow further response by direct PCB measurements downstream. To this end, turbidity monitors will be placed around each dredging or debris area undergoing remediation. Information from these monitors will provide continuous feedback to the operators, allowing adjustments in the operations to be made as needed in real time.

Objective XI: Determine the amount of suspended solids released by the remedial operations to provide an indication of PCB export.

1. Calculations presented in Attachment C indicate that the primary release mechanism of dredging-related contamination is resuspension of contaminated sediment. Thus, an increase in suspended solids should correlate with an increase of PCB

contamination. Samples will be collected for suspended solids analysis will be collected every three hours. At the near-field stations, these samples will be collected during the hours of operation. At the far-field stations, these samples will be collected on a 24-hours per day basis. These data, combined with the results of the far-field PCB analytical results, can be used to verify the relationship between suspended solids and PCB concentrations, and also provide a means of adjusting the suspended solids based resuspension criteria.

Objective XII: Verify that the NYSDEC surface water quality regulations are not violated during the remediation.

1. NYSDEC has water quality standards for pH and dissolved oxygen (DO). At both the near-field and far-field stations, pH and DO will be monitored discretely each time a sample is collected. These parameters plus conductivity will also provide a measure of quality assurance for the data collected.

Objective XIII: Determine a measurement that will be a real time indicator of PCB concentrations.

1. Because the primary release mechanism of dredging-related contamination during remediation is thought to be resuspension of contaminated sediment, a real time indication of suspended solids could provide a real time indication of PCB contamination in the water column. To develop the correlation between suspended solids and turbidity, continuous turbidity measurements will be required in the near field and the far field. Suspended solids will be measured every three hours. Literature reviews on this topic have shown that the correlations between suspended solids and turbidity are site-specific and can have large associated error. Particle counter measurements will be required to provide a second means of developing a continuous suspended solids measurement. Particle counter monitors will be required continuously at the four far-field stations from the TI Dam to Waterford. Discrete particle counter measurements will be required at the near-field stations.

Objective XIV: Verify the selection of the monitoring locations.

1. The locations of the far-field and near-field monitoring stations were selected based on several considerations, including near-field and far-field monitoring, ease of access, and level of planned dredging activities. The suspended solids and PCB analyses will be used to verify that these locations are appropriate. Monitoring of the far-field station less than one mile from the remediation will be required even though the PCB measurements will not be used for comparison to resuspension criteria during Phase 1. These results will determine if the station is heavily impacted by the nearby remediation and provide verification of the requirement that far-field stations be more than 1 mile from the remediation. (Monitoring for compliance with the far-field net suspended solids resuspension criteria will be required each day, no matter how close the remediation is to the far-field stations.)

4 Estimates of the Tolerable Error for the Monitoring Sampling Frequency Using Decision Error Feasibility Trials (DEFT) Software

4.1 Introduction

The EPA's guidance on data quality objectives (USEPA, 2000) was used in the development of the monitoring program for the Phase 1 dredging operation. This guidance describes a seven-step process for the identification of the decision points and data needs relating to the environmental problem to be addressed. With regard to PCB releases via resuspension during the Phase 1 operation, there are concerns to be resolved:

- How can EPA verify that PCB concentrations in the Upper Hudson River are in compliance with the resuspension criteria?

The focus of this analysis will be determining the appropriate sampling program, and specifically the frequency of sampling, that must be implemented to address these concerns.

In the following discussion, the DQO process (EPA QA-G4; EPA, 2000) is applied as outlined below:

1. State the Problem

2. Identify the Decision

3. Identify the Inputs to the Decision

4. Define the Boundaries of the Study

5. Develop a Decision Rule

6. Specify Tolerable Limits on Decision Errors

7. Optimize the Design for Obtaining Data

A separate discussion is provided for each question. A summary of the sampling requirements is provided is included in Section 1 of the Resuspension Performance Standard.

4.2 Development of Data Quality Objectives

4.2.1 Statement of the Problem

USEPA needs to verify that water column concentrations of PCBs in the Upper Hudson are below the Resuspension Standard criteria, thereby permitting unfettered dredging

operations. If PCB concentrations are not within acceptable levels, then additional monitoring and possible modifications to the engineering operations may be required.

USEPA staff represent the decision makers who will consult with GE, NYSDEC, water supply operators, local government representatives and non-government organizations.

The conceptual model is defined as follows:

> PCB loads and concentrations within the Upper Hudson are currently derived from sediment-based sources that contribute about 50 to 200 ng/L to the water column under typical flow conditions. These concentrations constitute baseline conditions. Dredging of contaminated sediments will add to this water column burden to some degree. Anticipated load additions due to dredging are expected to be less than 300 g/day (Evaluation Level threshold) under normal routine dredging for a 6-year remediation program. This is especially true for Phase 1 since the operation is planned at only half of the annual production rate anticipated in Phase 2.

> Although the mean daily Total PCB load increase due to dredging is expected to be well below 300 g/day, instantaneous conditions may result in momentary fluxes that are much higher. Consistent Total PCB loads higher than 300 g/day are considered indicative of problems in the dredging operation and warrant further study. Exceedance of 300 g/day does not constitute an immediate risk to human or ecological health but rather will delay the recovery of the river if allowed to continue for long periods of time. Similarly, exceedance of 600 g/day does not represent an immediate risk but again, loads at this level will delay the river's recovery if allowed to continue for an extended period of time.

> Total PCB concentrations in excess of 350 ng/L do not of themselves represent a risk to downstream users so long as levels remain below the drinking water maximum contaminant level (MCL) of 500 ng/L (total) PCBs. However, the proximity of this level (350 ng/L) to the MCL warrants more careful scrutiny and closer observation if 350 ng/L is exceeded due to the short transit time from the dredging area to the nearest public water supply intakes (two to seven days).

> Suspended solids data will provide an indication of increased PCB contamination in the water column. Net far-field suspended solids concentrations must be below 12 mg/L to be at routine levels and below 24 mg/L to be at or below the Evaluation Level. Net near-field suspended solids concentrations (as defined in the Resuspension Standard) must be below 60 mg/L, 100 mg/L or 700 mg/L depending on the location of the station relative to the dredge and the river section in which dredging is occurring. The duration of the exceedances provides an indication of the severity of the exceedance and the required response.

4.2.2   Identify the Decision

Depending on the magnitude of the dredging-related PCB load increase, USEPA may decide to do one or more of the following as described in Section 1 of this document:

- Increase monitoring frequency,
- Modify monitoring techniques,
- Modify dredging operations,
- Add additional engineering controls to the dredging operation, and
- Suspend the dredging operation until the PCB release problem has been resolved

For this decision, the primary question is "Are water column concentrations in compliance with the resuspension criteria?" If this is not true, required actions involve collection of additional samples to further define the PCB loads if the requirements of the first decision statement are met, with further increases in monitoring and the possibility or requirement of engineered modifications to the operation, as described in the standard.

4.2.3   Identify the Inputs to the Decision

To determine net PCB loads due to dredging (i.e., the total load less the baseline or historical load), the following data are needed:

a) Instantaneous and mean daily river flow at all monitoring locations

b) PCB concentrations at multiple monitoring locations, including the first far-field station downstream of the dredging operation and extending to Waterford.

c) PCB concentration at a location upstream of the dredging operation (specifically Rogers Island)

d) Suspended solids concentrations

e) Total organic carbon (TOC) on suspended solids

f) Dissolved organic carbon content (DOC; *i.e.,* TOC on filtered water samples)

g) Historical concentrations of PCBs, suspended solids, TOC on suspended solids at each of the main monitoring locations

Items a) through f) are used to characterize the actual conditions during dredging. Item g) provides a basis for comparison to establish the net load relative to the historical baseline conditions. The difference between baseline conditions and conditions measured during dredging is the net increase in PCB concentration due to dredging at each monitoring location. The product of mean daily flow and this concentration difference yields the estimate of the net load increase for comparison against the load-based criteria. Suspended solids and PCB concentration data will be used together to examine the usefulness of a suspended solids-PCB correlation to estimate PCB levels based on suspended solids monitoring alone.

The methods for sample analysis include:

1. PCB congeners with a detection limit of 0.5 ng/L total PCBs. The effective congener detection limit is roughly 0.05 ng/L Currently this can only be achieved by one of the following: EPA's dual column GC/ECD method, Standard Method 1668A or GE's modified Green Bay Method.
2. Total Suspended Sediment with a detection limit of 0.1 mg/L, by Analytical Method ASTM D3977-97, Standard Test Method for Determining Sediment Concentration in Water Samples, or equivalent. No subsampling of a sampling container is permitted.
3. Organic carbon on the suspended solids can be done via a Total Organic Carbon method or by a combustion technique but must be sensitive down to 0.1% (1000 mg/kg) on the suspended solids.
4. Dissolved organic carbon method should have a detection limit of 0.5 mg/L, such as ASTM Method D4839-03 [0.1 mg/L] or EPA 415.2 [.05 mg/L].

4.2.4   Define the Boundaries of the Study

The boundaries of the site are defined as the shorelines of the Hudson River, excluding its tributaries, between the Fennimore Bridge at Hudson Falls and the Federal Dam at Troy. The Fennimore Bridge is included as the upper boundary, rather than the northern end of Rogers Island, because of the potential for PCB releases associated with the remediation of the GE Hudson Falls facility that will be taking place at the same time or just prior to the sediment remediation.

In recognition of the need to simplify monitoring, monitoring locations will be chosen considering ease of access as well as project data needs. The ease of access consideration leads to the selection of the Fennimore Bridge, Rogers Island, Schuylerville, Stillwater and Waterford locations, which are all accessible by bridge. These locations also roughly divide the river into 10 to 15 mile segments, providing sufficient resolution to identify potential PCB sources by location. The separation of these locations also reflects the desire to allow natural hydrodynamic processes to homogenize PCB concentrations in the river, simplifying the sampling process to some degree.

Given that most of the dredging is scheduled for the TI Pool, an additional monitoring location is identified at the TI Dam so as to better identify loads originating in this reach.

Because dredging-related releases will depend on many factors related to dredge operation, sediment type and location with in the river, the PCB load is expected to vary significantly over time. Daily monitoring is considered a minimum basis for determining compliance with the lowest (most stringent) secondary criterion of 300 g/day. Higher frequency monitoring is needed to document and understand the sources of PCBs to the water column when this threshold is exceeded.

The loads released by dredging are expected to vary rapidly over time and thus will need to be reviewed daily. Sampling when routine conditions are expected will measure the daily variability. However, the weekly condition as defined by a 7-day running mean

calculated daily will be used to test compliance with the load-based criteria. In this manner, compliance with the long-term load criterion can be confirmed while also collecting data to demonstrate that more significant exceedances of PCB concentration criteria (*e.g.,* exceeding 350 or 500 ng/L) have not occurred.

The transit time of water from the TI Pool to Waterford is expected to vary from two to seven days, depending inversely on flow. As a result of the normal dispersion and settling processes, the intensity of any short-term PCB release is expected to be diminished as the river travels from TI Pool to Waterford. Thus, for a dredging operation in the TI Pool, the discrete sample collected at TI Dam effectively integrates a much shorter time period than a sample obtained at Waterford. Thus collecting samples along the Upper Hudson serves to examine both short-term (one hour duration) and longer-term (one- to two-day duration) PCB loads and PCB concentrations. Both measures are needed to assess the success of the resuspension controls.

The sampling program must reflect the need to respond to gradual increases in long-term impacts such as PCB mass transported as well as to consideration of acute PCB concentrations at downstream public water supplies. For this reason, both long-term averages (7-day and 4-week periods) and daily results are needed. To address the protection of downstream water supplies, 24-hour turn-around times are needed for the two monitoring stations downstream of, but closest to, the dredge operation. For Phase 1, these are expected to be the TI Dam and Schuylerville stations. From these considerations and those of the standard itself, the decision units are the loads as measured weekly to monthly (4 week) and the concentrations measured daily.

For locations farther downstream, the results from the two far-field stations closest to the dredging operations provide some confidence that levels farther downstream will be acceptable (or at least that major excursions will be known). However, due to the highly variable nature of the PCB release process, samples must still be collected from locations farther downstream and levels confirmed to be in compliance with the standard. These samples can have a longer turn around time, on the order of 7 days from collection to result, since their role is primarily confirmational. These samples are necessary during Phase 1 but may be dropped in Phase 2, depending on the success of the suspended solids monitoring and the actual PCB loss rates.

4.2.5   Develop a Decision Rule

The decision rules are derived from the performance standard criteria described in Section 1 and justified in Sections 2 and 3 of the main document. The decision rule is designed to test compliance with the standard criteria.

> *The arithmetic mean is selected as the primary measure since it reflects an integration of several measures and representative of the integrated PCB load over the averaging period. Compliance with each of the resuspension criteria is the primary focus of this DQO discussion.*

4.2.6 Specify Tolerable Limits on Decision Errors

Current estimates of PCB release due to dredging, as developed in other attachments to this document, indicate that PCB loads and concentrations are likely to fall below the action level criteria during most of the operation. That is, when viewed on a daily or weekly basis, momentary flux variations will average out so as to fall below the action level criteria. Additionally, the threshold criteria developed for the decision rules do not represent conditions immediately dangerous to human health or the environment. Based on this, the null hypothesis for the decision rule is taken as the condition that the river is in compliance (*i.e.,* the river flux or concentration of total PCBs is below the criteria value). This approach also takes into consideration the fact that monitoring will continue and that confirmation of any day's decision about dredging releases and water column concentration is only 24 hours away, resulting from the next day's sampling.

USEPA's Decision Error Feasibility Trials Software (DEFT (USEPA, 2001)) was used to develop the sampling requirements for this program. The results of this analysis are presented in Table 1. As defined in USEPA (2001):

> A *false acceptance* decision error occurs when the sample data lead you to decide that the baseline condition is probably true when it is really false.

> A *false rejection* decision error occurs when the limited amount of sample data lead you to decide that the baseline condition is probably false when it is really true.

> The *gray region* is a range of true parameter values within the alternative condition near the Action Level where it is "too close to call."

False acceptances were minimized because this is the more serious error. In general, decisions that are more critical, such as confirmation of exceedance of the Resuspension Standard which requires shut down or exceedance of the Control Level which requires intense monitoring and implementation of engineering solutions, require a large number of samples and have greater certainty than the less critical decisions. For the suspended solids measurements, it is clear that the implementation of a continuous monitor capable of estimating suspended solids concentrations will be needed to having a reasonable amount of certainty in these decisions. The low level of certainty is tolerable only because any decisions made as a result of exceedance of the suspended solids will be confirmed by measurements of PCB concentrations in the impacted water column.

For PCB measurement-based decisions, a false acceptance rate of 5 percent or less was sought, with lower rates sought when an incorrect decision would yield an unnecessary halting of the operation or an engineering improvement. The rate of 5 percent was selected as an acceptable error for the lower action level criteria since exceedance of the action level criteria only initially induces additional monitoring which will quickly confirm the exceedance. This error rte reflects a balance between by the desire to keep monitoring requirements as low as possible while still providing protection.

4.2.7 Optimize the Design for Obtaining Data: Results of the Analysis

The final sampling requirements for the standard were developed using DEFT (USEPA, 2001), a program to estimate sampling requirements based on a project-specific error rate. The analysis of the various criteria and acceptable *gray region* around each criterion yielded the results given in Table G-2. For all criteria except the confirmation of the 500 ng/L exceedance, the null hypothesis assumed that river conditions were in compliance. Table G-1 summarizes the various criteria, the associated gray region, the sampling frequency required by the resuspension standard and the false acceptance and false rejection levels. The table is organized by measurement type (*i.e.,* PCB and suspended solids).

Two important assumptions were made to develop the error rate values in the table. Specifically, there is no site-specific data on the expected variance of water column conditions related to dredging. As a result, the extensive analysis of variance compiled in Attachment A was used to this end. A nominal coefficient of variance was assumed for PCB and for suspended solids based on the variance observed under baseline conditions. For PCB measurements (both Total PCBs and Tri+), the coefficient of variance is assumed to be 25 percent. For suspended solids, the coefficient of variance was assumed to be 75 percent.

This section also includes a set of figures illustrating the statistical calculations used to estimate the error rates. Figures 1 to 26 represent the calculations for each line in Table G-1.

The table shows that the higher level of sampling associated with the higher action levels and the and Resuspension Standard yield low false error rates, as expected, reflecting the need to be accurate before taking costly actions or improvements. In some instances, the false rejection rate is fairly high, indicating that additional sampling may be unnecessarily triggered. However, this represents a protective approach from the perspective of the safety of the public water supplies. Additionally, the higher monitoring rates will quickly confirm the need to remain at the action level thought to be exceeded.

Higher error rates are estimated in transitions from routine conditions to the Evaluation and Concern Levels, reflecting the relative low sampling rate required for routine sampling. Also shown in the table is the one week confirmation result (*i.e.,* the error rate for the combination of one week of routine monitoring and one week at the action level). In each instance the false acceptance error is brought below 5 percent, thereby confirming the need to sample at the higher rate or indicating that sampling at the routine rate may be resumed.

The results for monitoring requirements for exceedance of the standard demonstrate the need for the intensive sampling specified. In this instance the river is assumed be in exceedance of the standard. Four additional discrete samples (Figure 10) do not provide sufficient certainty given that the next day's decision will involve the temporary halting

of the dredging operations, a costly choice. However, by collecting hourly composites, the power of the same four analyses is greatly improved and the 5 percent false acceptance rate is attained.

The table also shows the results for the long-term integrative samples. These samples will serve to confirm the results of daily routine monitoring or show the need for more frequent sampling. The results assume the automated collection of eight samples per day over a one- to two-week period.

The results for suspended solids illustrate the need to use a continuous sampling system such as a turbidity probe. In the lower portion of the table, results for the discrete sampling program are compared with those that can be achieved with a continuous probe recorded once every 15 minutes. In almost all cases, the continuous reading probe provides more than an order of magnitude improvement in the expected error rate. Better rates can be achieved wit the continuous probes by simply recording more frequently. Note that this analysis does not consider any uncertainty introduce by use of a probe over discrete samples. Nonetheless, given a semiquantitatve relationship between the probe and actual suspended solids levels, it is highly likely that the probes will provide a substantial reduction in the expected error rates for suspended solids monitoring, thereby reducing unnecessary additional PCB sampling prompted by a false indication.

5   References:

USEPA, 2000. Guidance for the Data Quality Objectives Process EPA/600/R-96/055. August 2000.

USEPA, 2001. Data Quality Objectives Decision Error Feasibility Trials Software (DEFT) - USER'S GUIDE. EPA/240/B-01/007. September 2001.

Tables

Table G-1
Summary of Sampling Frequency Requirements and Expected Error Rates

Analysis	Transition	Detail	Sampling Time Period	Action Level	Number of Samples	Grey Region Limit	False Rejection Error Limit - α (%)	False Acceptance Error Limit - β (%)	Figure Number
Total PCB Sampling Requirements (25% CV)									
	Routine to Evaluation Level	Routine to > 300 g/day	1 week	300 g/day	7	400 g/day	7.5	5	1
	Routine to Concern Level	Routine to > 600 g/day	1 week	600 g/day	7	700 g/day	25	15	2
	Confirmation of the Concern Level	Confirmation of > 600 g/day	1 week routine + 1 week	600 g/day	28	700 g/day	5	4	3
	Routine to Concern Level	Routine to > 350 ng/L	1 week	350 ng/L	7	400 ng/L	28	20	4
	Confirmation of the Concern Level	Confirmation of > 350 ng/L	1 week routine + 1 week	350 ng/L	28	400 ng/L	10	5	5
	Evaluation to Concern Level	300 g/day to > 600 g/day	1 week routine + 1 week	600 g/day	21	700 g/day	10	5	6
	Concern to Control Level	600 g/day-7-day to 4-week average	1 week routine + 3 weeks	600 g/day	70	700 g/day	0.5	0.2	7
		350 ng/L - 7-day to 4-week average	1 week routine + 3 weeks	350 ng/L	70	400 ng/L	1	0.9	8
	Control Level to Resuspension Standard Threshold	350 ng/L to >500 ng/L	2 days	500 ng/L	8 composites of 6 aliquots each	550 ng/L	16.5	5	9
	Resuspension Standard Threshold	Confirmation of > 500 ng/L	1 day routine + 1 day	500 ng/L	5	400 ng/L	30	15	10
		Confirmation of > 500 ng/L (24 hours)	1 day	500 ng/L	4 composites of 6 aliquots each	400 ng/L	7	5	11
	Routine to Concern Level or Concern to Control Level	Continuous Total PCB 1-week or 2-week deployment	1 week or 2 weeks	350 ng/L	2 composites of 56 aliquots each	400 ng/L	6.5	5	12
Suspended Solids Sampling Requirements (75% CV)									
	Routine to Evaluation Level	Far-field - Baseline to > 12 mg/L	1 day (3 hrs for 24 hrs)	14 mg/L	8 (discrete)	21 mg/L	28	12.5	13
			1 day (15 min for 24 hrs)	14 mg/L	96 (continuous)	21 mg/L	0.1	0.1	14
	Routine to Concern Level	Far-field - Baseline to > 24 mg/L	1 day (3 hrs for 24 hrs)	26 mg/L	8 (discrete)	39 mg/L	28	12.5	15
			1 day (15 min for 24 hrs)	26 mg/L	96 (continuous)	39 mg/L	0.1	0.1	16
	Routine to Concern Level	Near Field - River Sections 1 and 3 Baseline to > 100 mg/L	6 hours (1 sample per 3 hours)	100 mg/L	3 (discrete)	150 mg/L	35	25	17
			6 hours (1 sample per 15 min)	100 mg/L	24 (continuous)	150 mg/L	6.6	5	18
	Routine to Concern Level	Near Field - River Section 2 Baseline to > 60 mg/L	6 hours (1 sample per 3 hours)	60 mg/L	3 (discrete)	90 mg/L	35	25	19
			6 hours (1 sample per 15 min)	60 mg/L	24 (continuous)	90 mg/L	6.6	5	20
	Concern to Control Level	Near Field - River Sections 1 and 3 Baseline to > 100 mg/L	1 day (3 hrs for 15 hrs)	100 mg/L	5 (discrete)	150 mg/L	28	20	21
			1 day (15 min for 15 hrs)	100 mg/L	60 (continuous)	150 mg/L	0.7	0.5	22
	Concern to Control Level	Near Field - River Section 2 Baseline to > 60 mg/L	1 day (3 hrs for 15 hrs)	60 mg/L	5 (discrete)	90 mg/L	28	20	23
			1 day (15 min for 15 hrs)	60 mg/L	60 (continuous)	90 mg/L	0.7	0.5	24
	Routine to Evaluation Level	Near Field Baseline to > 700 mg/L	3 hours (1 sample per 3 hours)	700 mg/L	2 (discrete)	1000 mg/L	40	30	25
			3 hours (1 sample per 5 min)	700 mg/L	36 (continuous)	1000 mg/L	16.5	5	26

Figures

Figure 1
Routine to Evaluation Level
Action level of 300 g/day

Estimated Performance Curve

Probability of deciding that the true mean is greater than the action level

1.0		0.050
0.9		
0.8	Action Level -->	
0.7		
0.6		
0.5	FR	FA
0.4		
0.3	Ho: mean ≤ 300	
0.2		
0.1		0.075
0.0		

0.0 70.0 140.0 210.0 280.0 350.0 420.0 490.0 560.0 630.0 700.0

True Mean Concentration

Simple Random Sampling
Action Level = 300.000
Cost = $2100.00
Sample Size = 7

Decision Error Limits
concentration prob(E) type
300.000   0.075   FR
400.000   0.050   FA

Figure 2
Routine to Concern Level
Action Level of 600 g/day
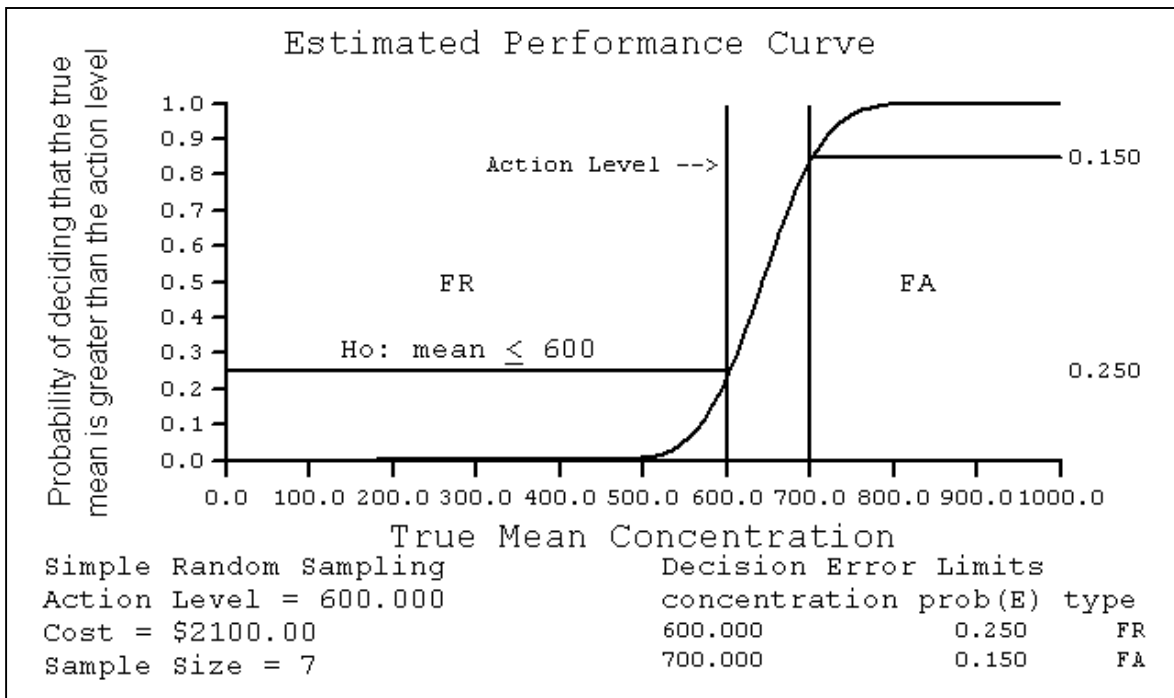
Estimated Performance Curve

Probability of deciding that the true mean is greater than the action level

1.0		
0.9		0.150
0.8	Action Level -->	
0.7		
0.6		
0.5	FR	FA
0.4		
0.3	Ho: mean ≤ 600	0.250
0.2		
0.1		
0.0		

0.0 100.0 200.0 300.0 400.0 500.0 600.0 700.0 800.0 900.0 1000.0

True Mean Concentration

Simple Random Sampling
Action Level = 600.000
Cost = $2100.00
Sample Size = 7

Decision Error Limits
concentration prob(E) type
600.000   0.250   FR
700.000   0.150   FA

Note: Figures generated from DQO – DEFT using a coefficient of variation for all total PCB cases of 25 percent.

Figure 3
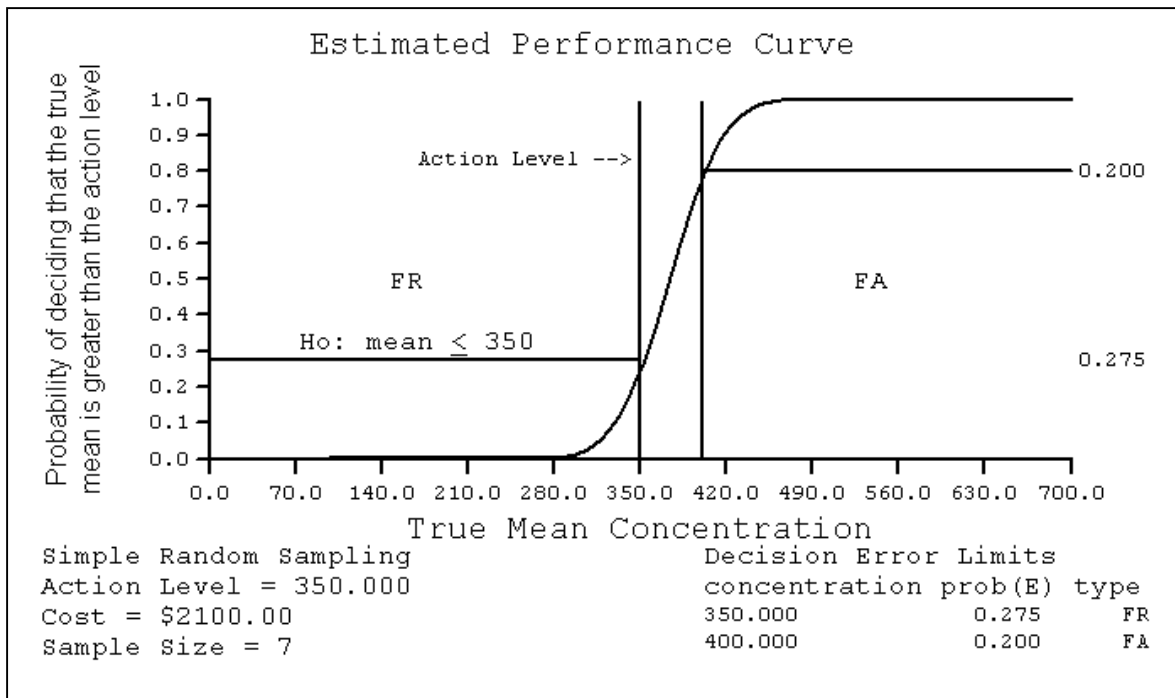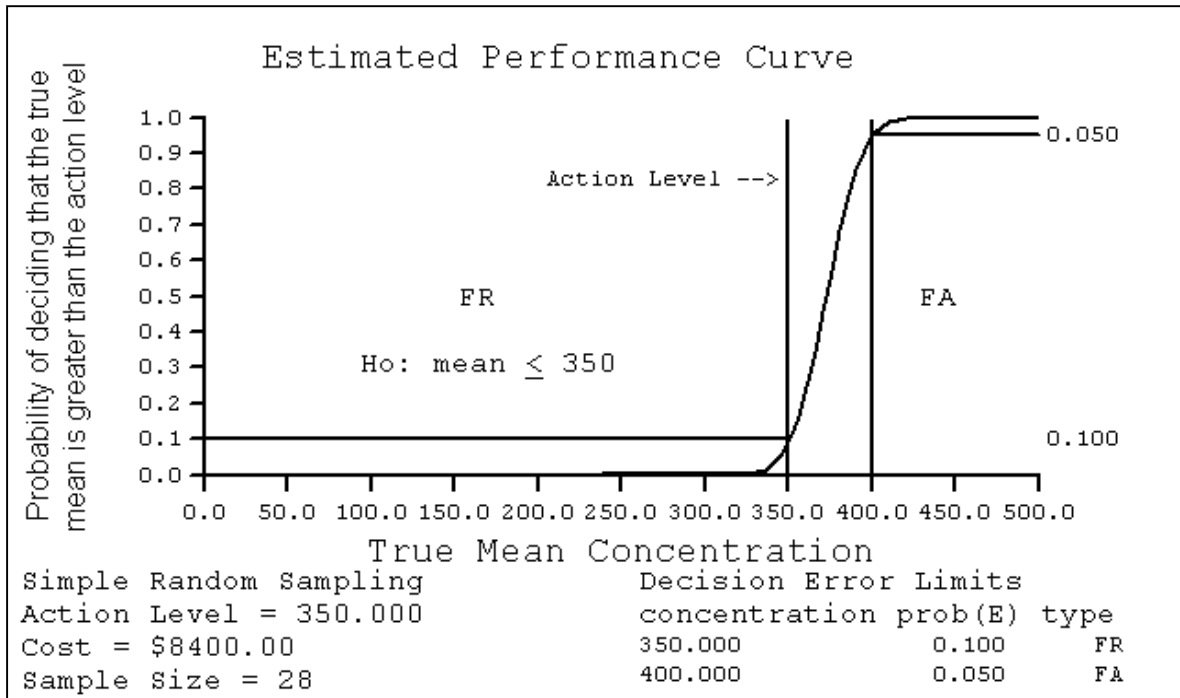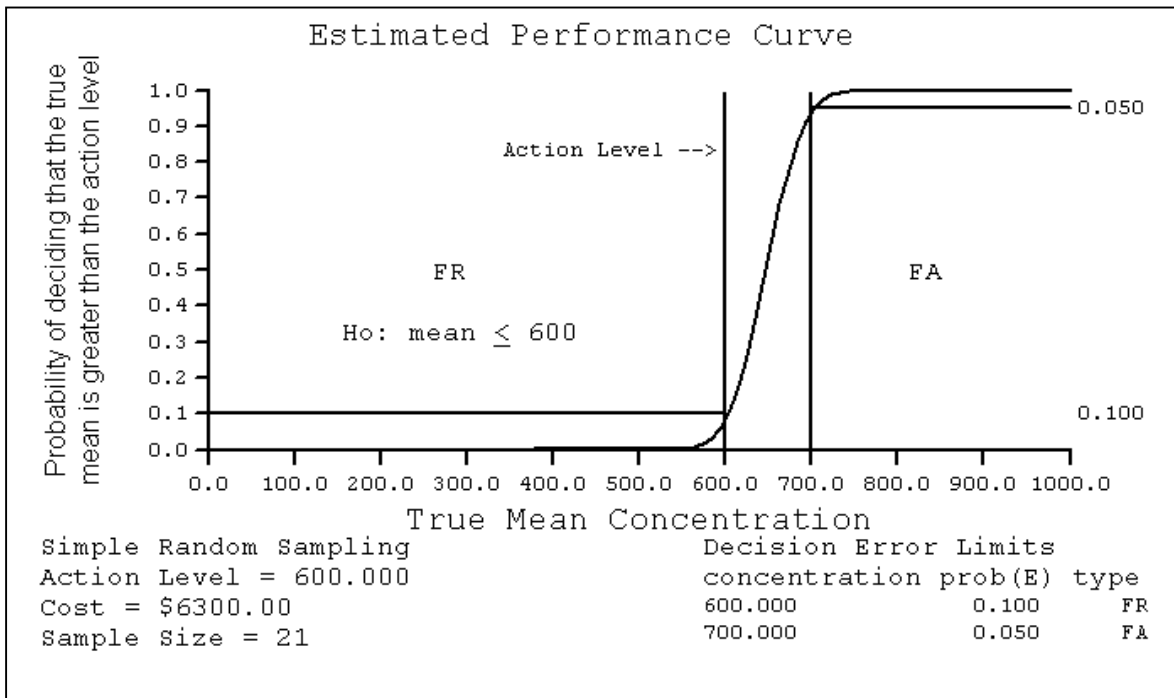**Confirmation of the 600 g/day**
Action Level of 600 g/day

Estimated Performance Curve

Probability of deciding that the true
mean is greater than the action level

Action Level -->

FR   FA

Ho: mean ≤ 600

True Mean Concentration

Simple Random Sampling   Decision Error Limits
Action Level = 600.000   concentration prob(E) type
Cost = $8400.00   600.000   0.050   FR
Sample Size = 28   700.000   0.040   FA
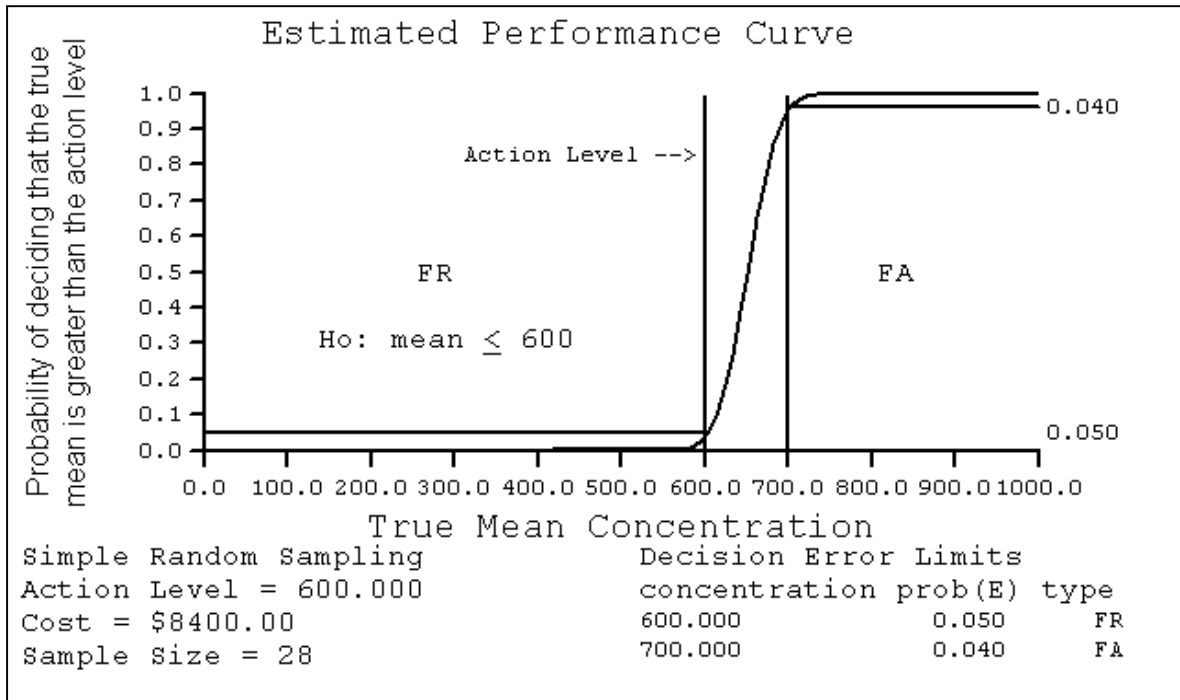
Figure 4
Routine to Concern Level
Action Level of 350 ng/L

Estimated Performance Curve

Probability of deciding that the true
mean is greater than the action level

Action Level -->

FR   FA

Ho: mean ≤ 350

True Mean Concentration

Simple Random Sampling   Decision Error Limits
Action Level = 350.000   concentration prob(E) type
Cost = $2100.00   350.000   0.275   FR
Sample Size = 7   400.000   0.200   FA

Note: Figures generated from DQO – DEFT using a coefficient of variation for all total
PCB cases of 25 percent.

Figure 5
Confirmation of the 350 ng/L
Action Level of 350 ng/L

Estimated Performance Curve

Probability of deciding that the true mean is greater than the action level

Action Level -->

FR

Ho: mean ≤ 350

FA

0.050

0.100

True Mean Concentration

Simple Random Sampling
Action Level = 350.000
Cost = $8400.00
Sample Size = 28

Decision Error Limits
concentration   prob(E)   type
350.000   0.100   FR
400.000   0.050   FA

Figure 6
Evaluation Level to Concern Level
300 g/day to 600 g/day

Estimated Performance Curve

Probability of deciding that the true mean is greater than the action level

Action Level -->

FR

Ho: mean ≤ 600

FA

0.050

0.100

True Mean Concentration

Simple Random Sampling
Action Level = 600.000
Cost = $6300.00
Sample Size = 21

Decision Error Limits
concentration   prob(E)   type
600.000   0.100   FR
700.000   0.050   FA

Note: Figures generated from DQO – DEFT using a coefficient of variation for all total PCB cases of 25 percent.

Figure 7
Concern Level to Control Level
Action Level of 600 g/day (7-day average to 4-week average)
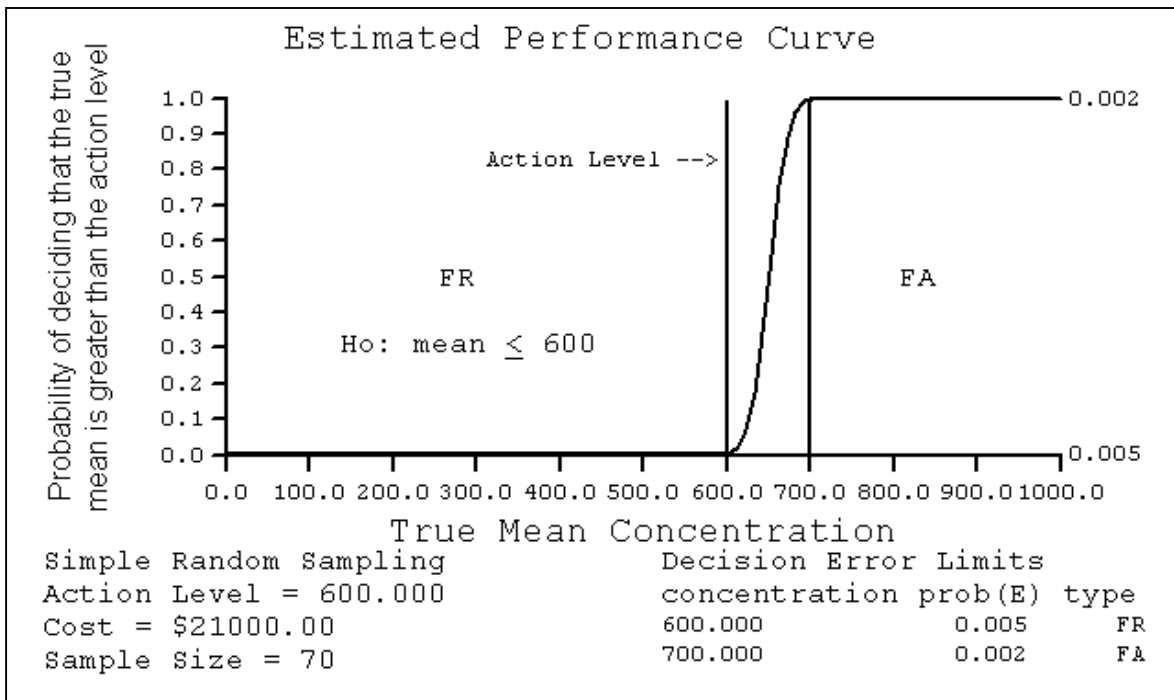
Figure 8
Concern Level to Control Level
Action Level of 350 ng/L (7-day average to 4-week average)
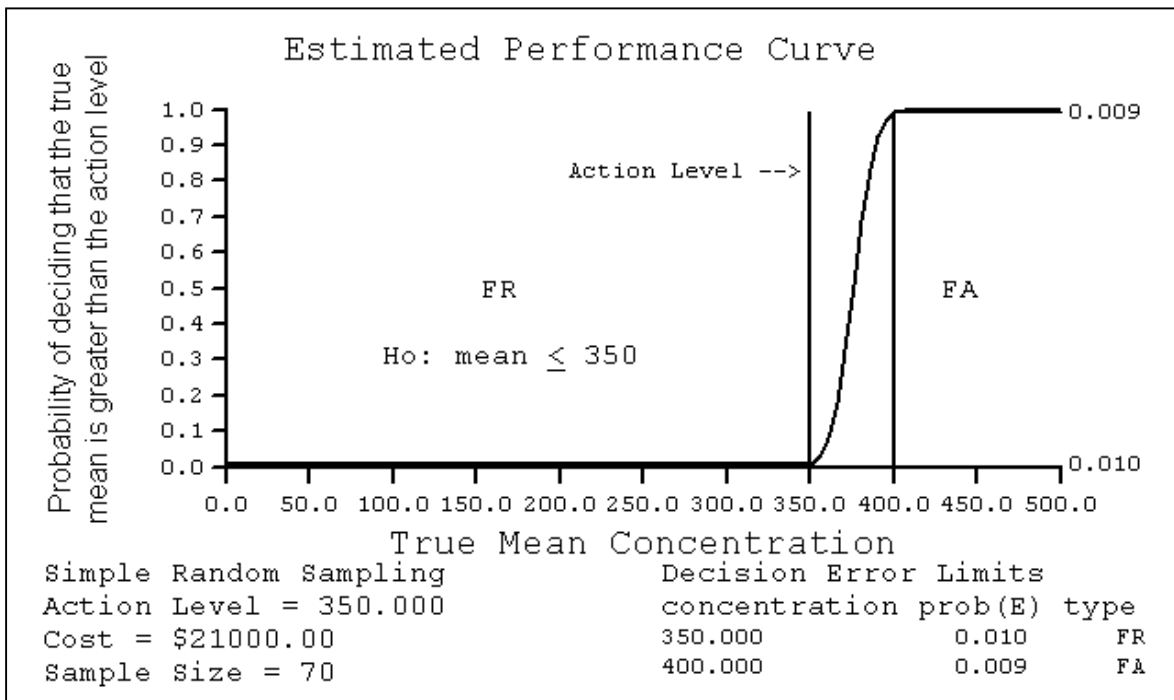
Note: Figures generated from DQO – DEFT using a coefficient of variation for all total PCB cases of 25 percent.

Figure 9
Control Level to Resuspension Standard Threshold
Action Level of 500 ng/L (2 days sampling; 4 sample/day; 6 aliquots/sample)
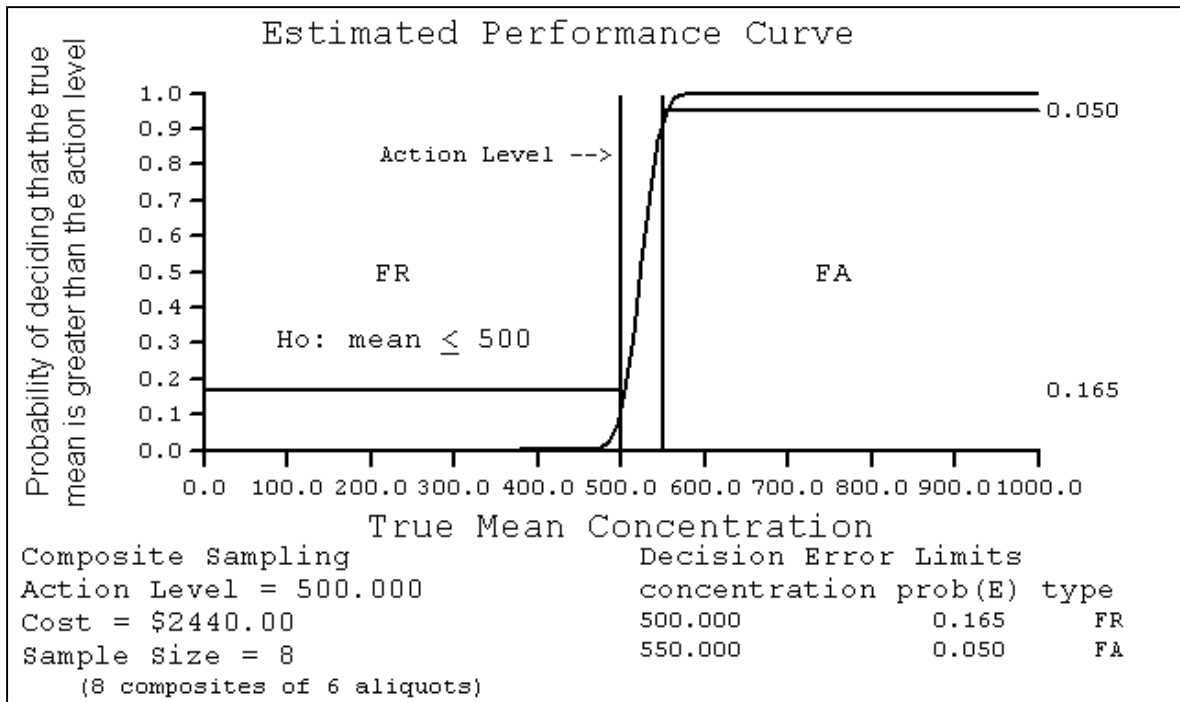
Figure 10
Resuspension Threshold
Confirmation of 500 ng/L
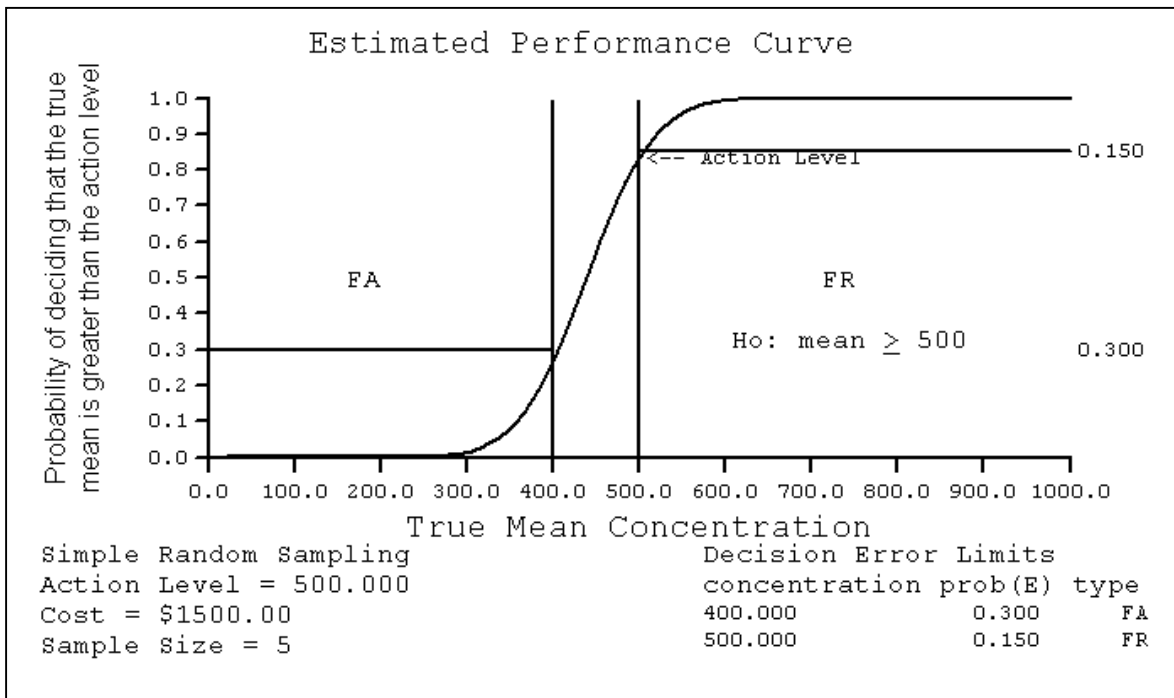
Note: Figures generated from DQO – DEFT using a coefficient of variation for all total
PCB cases of 25 percent.

Figure 11
Resuspension Threshold
Confirmation of 500 ng/L (24 hours; 4 samples of 6 aliquots)

Estimated Performance Curve

Composite Sampling
Action Level = 500.000
Cost = $1220.00
Sample Size = 4
    (4 composites of 6 aliquots)

Decision Error Limits
concentration prob(E) type
400.000   0.070   FA
500.000   0.050   FR
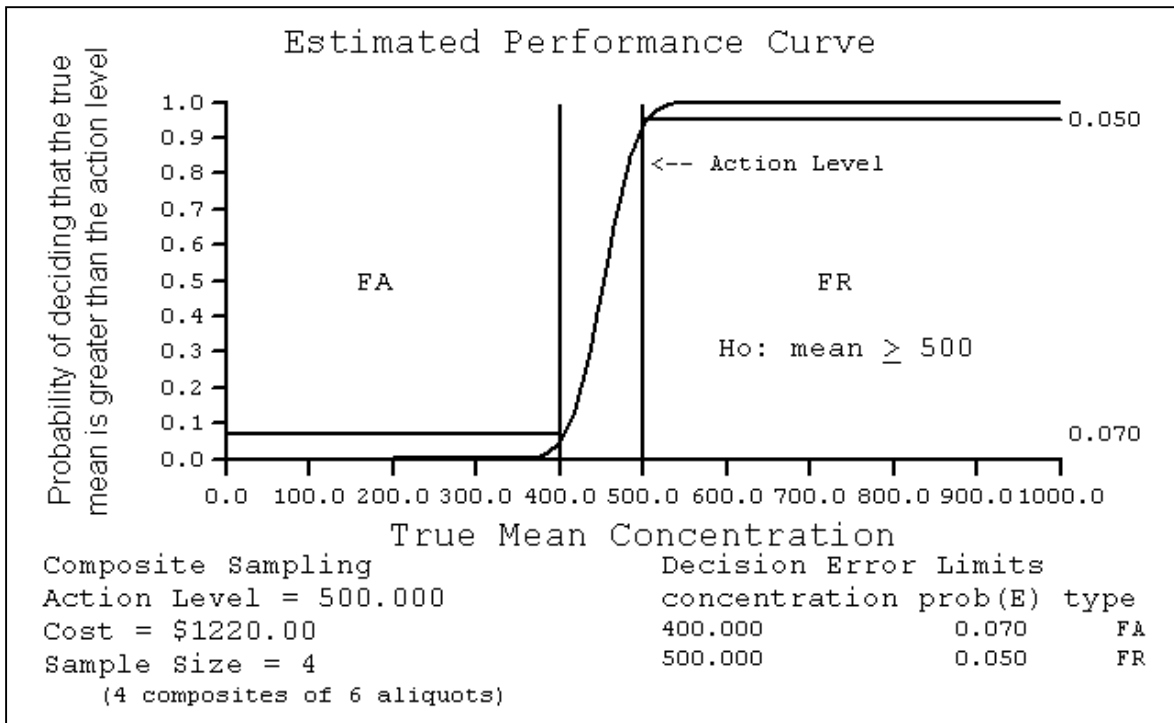
Figure 12
Routine to Concern Level (350 ng/L, 2-week deployment) or
Concern Level to Control Level (350 ng/L, 1-week deployment)
Continuous total PCB sampling requirements
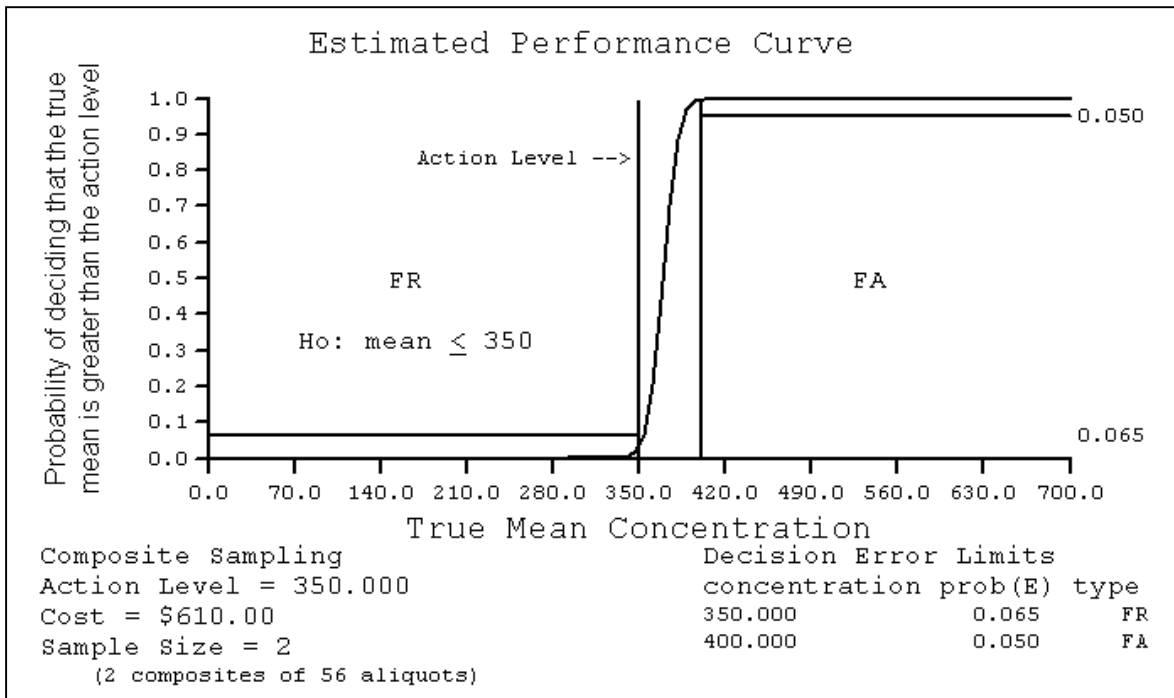
Estimated Performance Curve

Composite Sampling
Action Level = 350.000
Cost = $610.00
Sample Size = 2
    (2 composites of 56 aliquots)

Decision Error Limits
concentration prob(E) type
350.000   0.065   FR
400.000   0.050   FA

Note: Figures generated from DQO – DEFT using a coefficient of variation for all total PCB cases of 25 percent.

Figure 13
Routine to Evaluation Level
(Far-field Baseline to >12 mg/L with discrete samples every 3 hrs for 24 hrs)
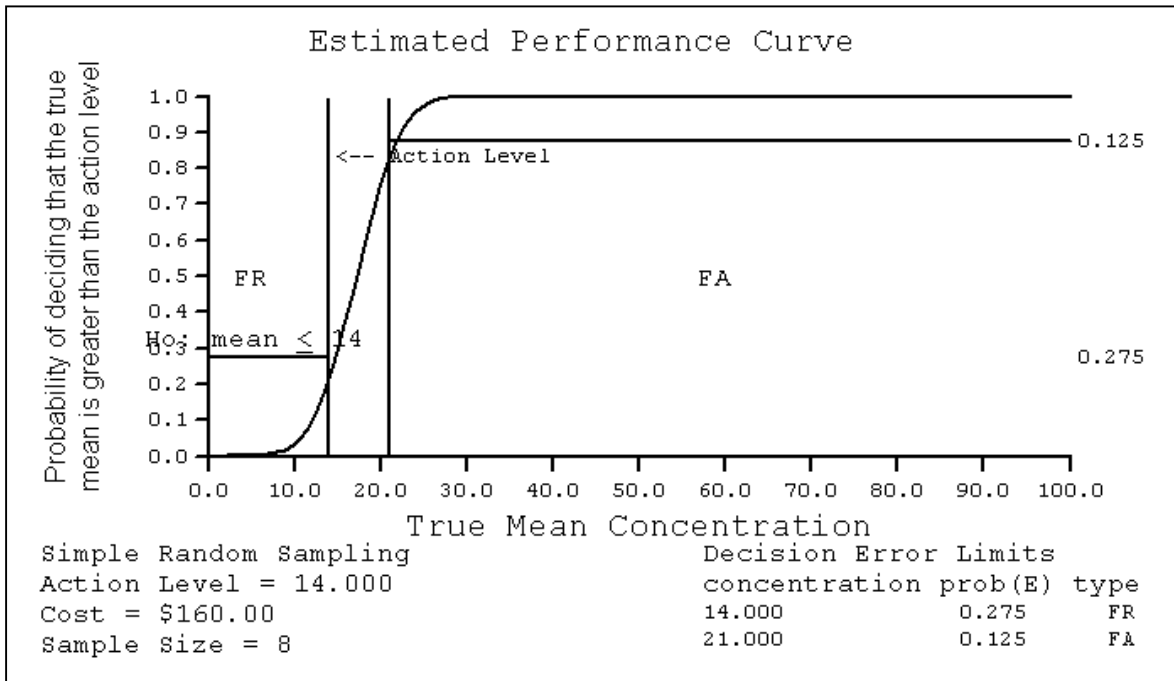
Figure 14
Routine to Evaluation Level
(Far-field baseline to >12 mg/L with continuous sampling every 15 min for 24 hrs)
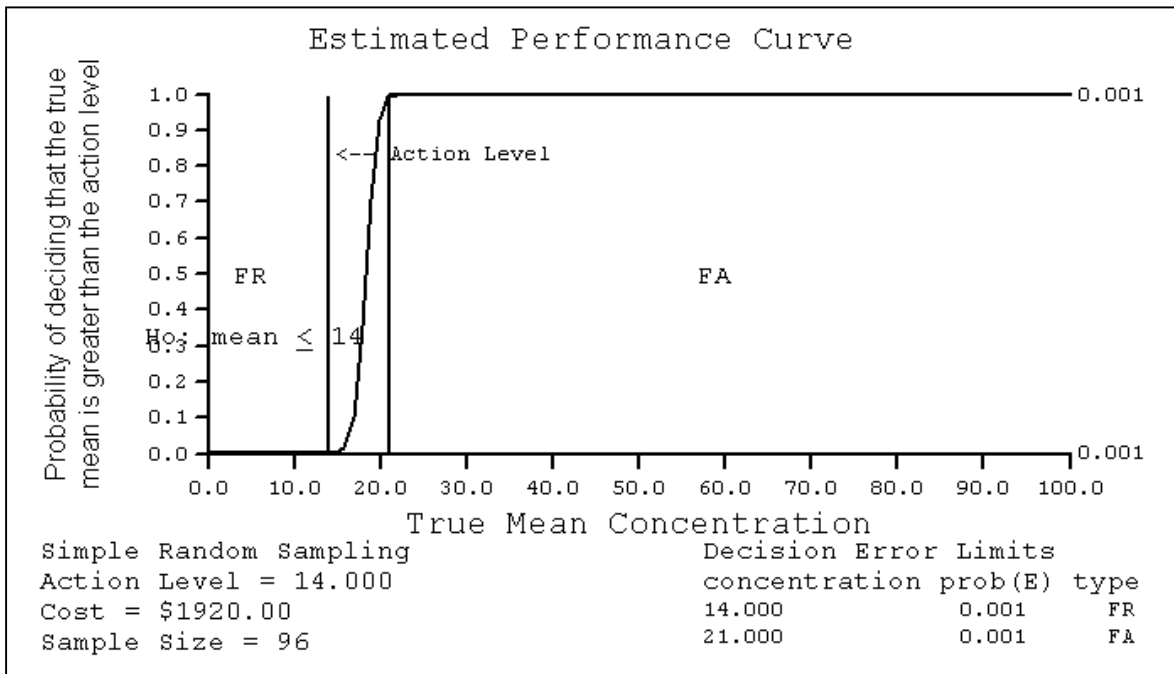
Note: The analysis is based on a baseline of Schuylerville conditions (Average TSS concentration from May-Nov of 2.4 mg/L with an average standard deviation from May-Nov of 1.87 mg/L) and coefficient of variation equal to 75 percent.

Figure 15
Routine to Concern Level
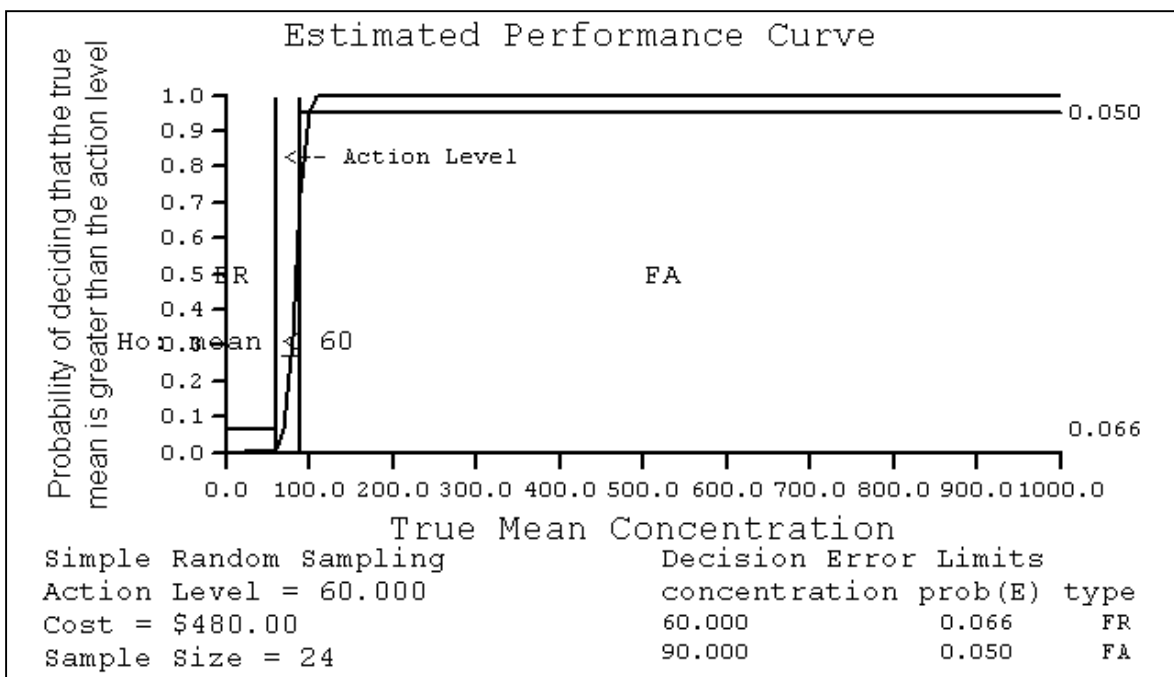**(Far-field Baseline to >24 mg/L with discrete samples every 3 hrs for 24 hrs)**
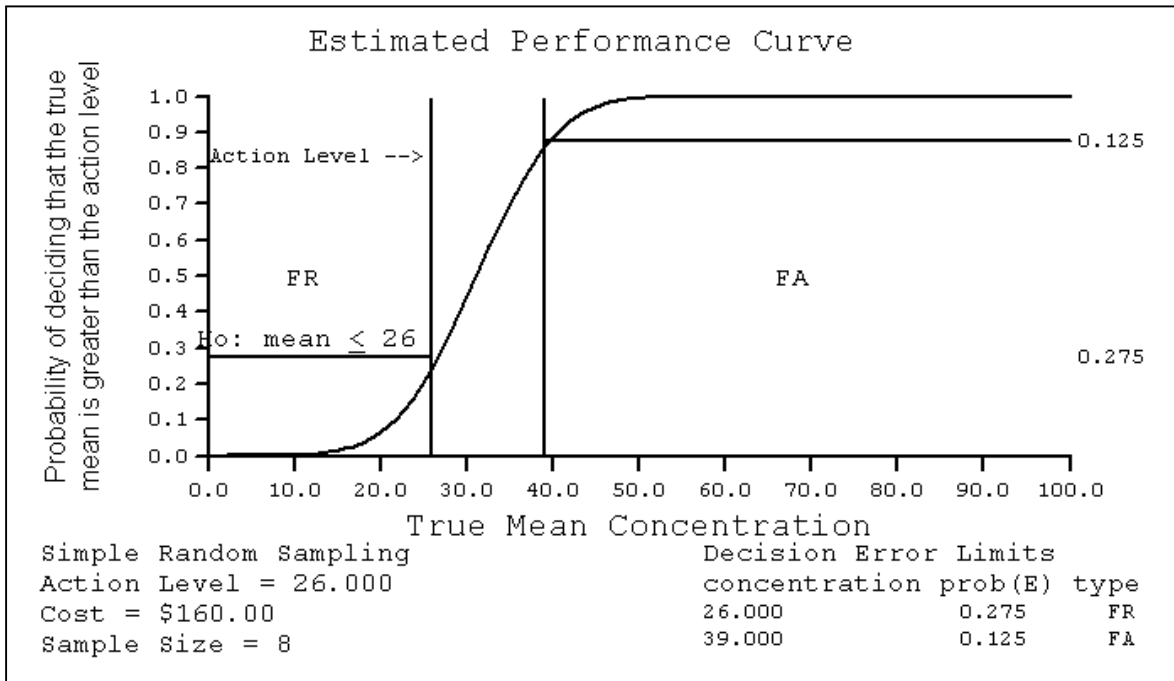
Figure 16
Routine to Concern Level
(Far-field baseline to >24 mg/L with continuous sampling every 15 min for 24 hrs)
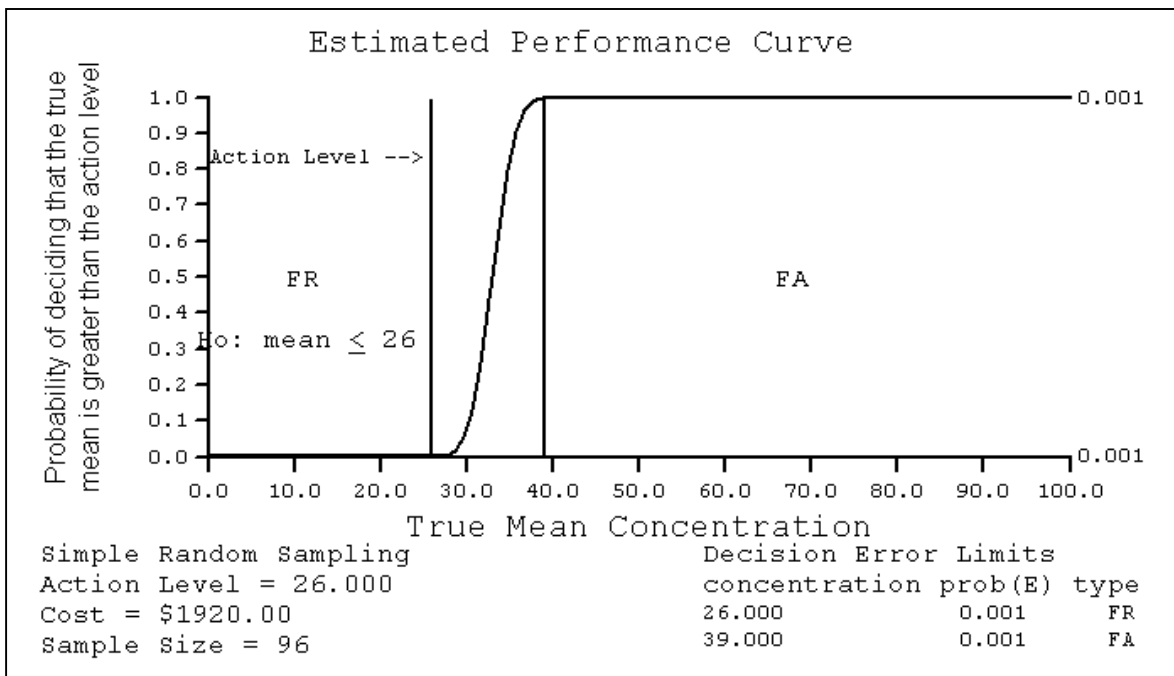
Note: The analysis is based on a baseline of Schuylerville conditions (Average TSS concentration from May-Nov of 2.4 mg/L with an average standard deviation from May-Nov of 1.87 mg/L) and coefficient of variation equal to 75 percent.

Figure 17
Routine to Concern Level Near-field River Sections 1 and 3
(baseline to >100 mg/L with discrete samples every 3 hours for 6 hours)
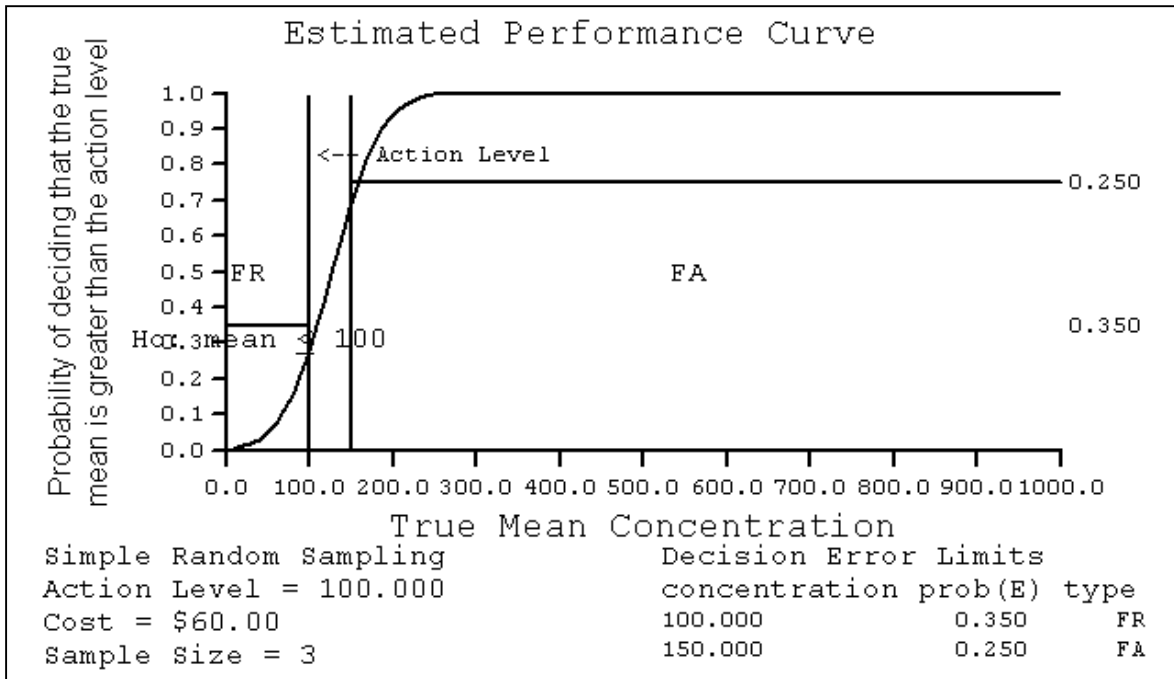
Figure 18
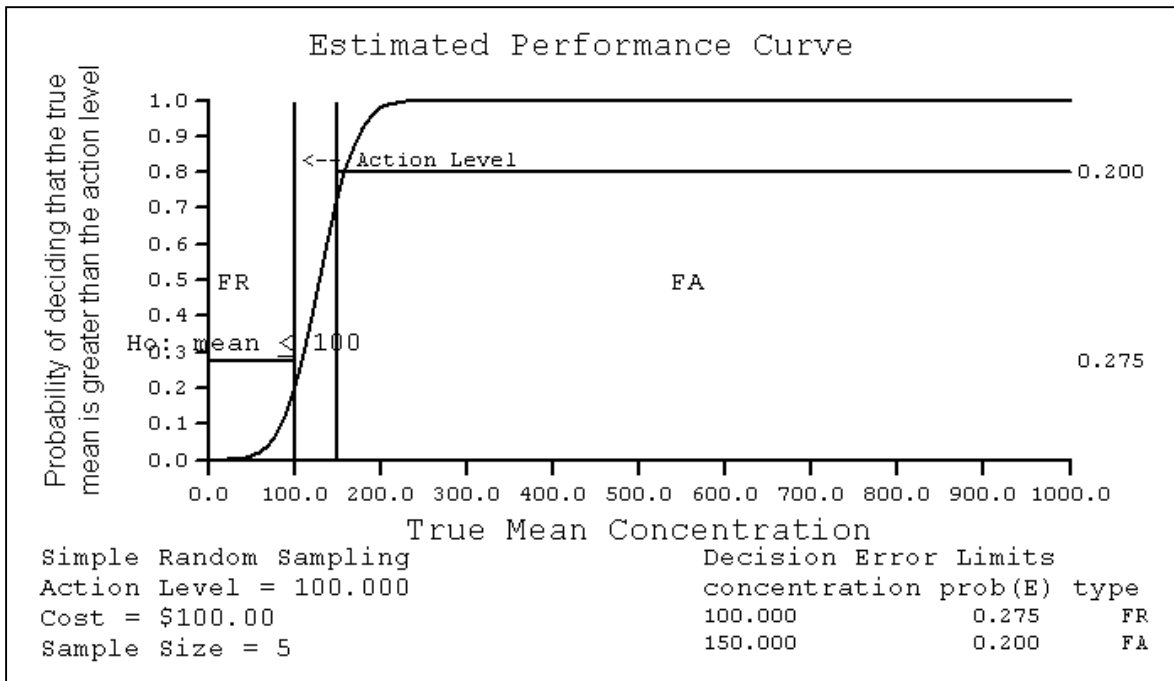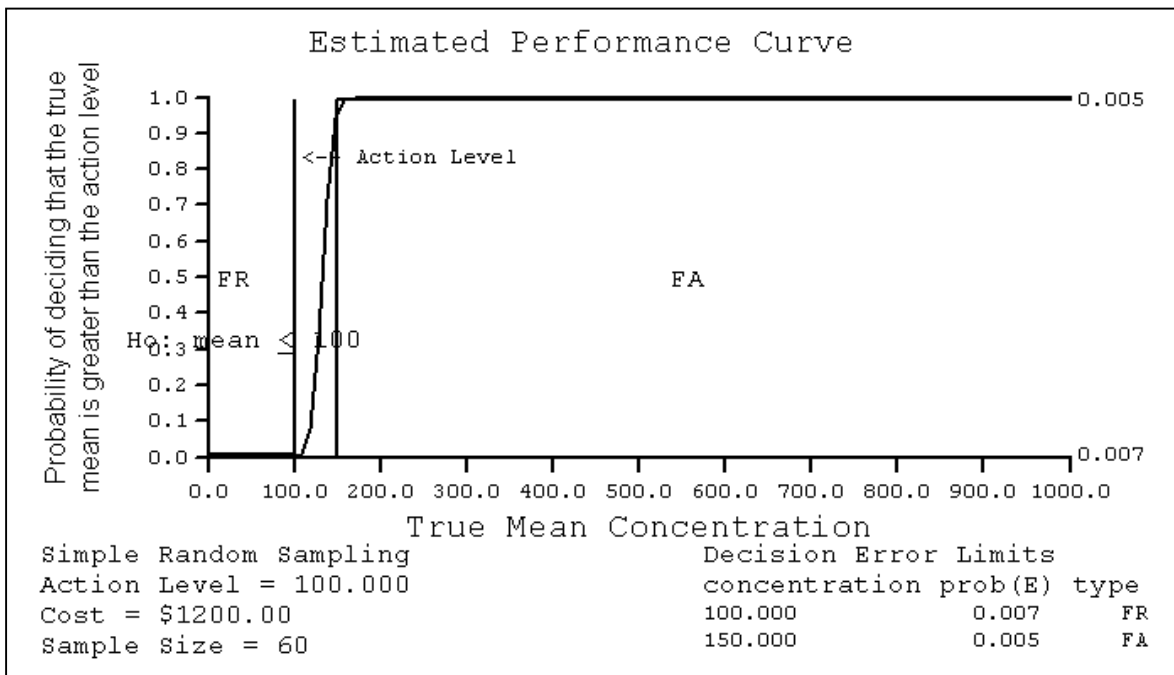**Routine to Concern Level Near-field River Sections 1 and 3**
(baseline to >100 mg/L with continuous sampling every 15 min for 6 hrs)
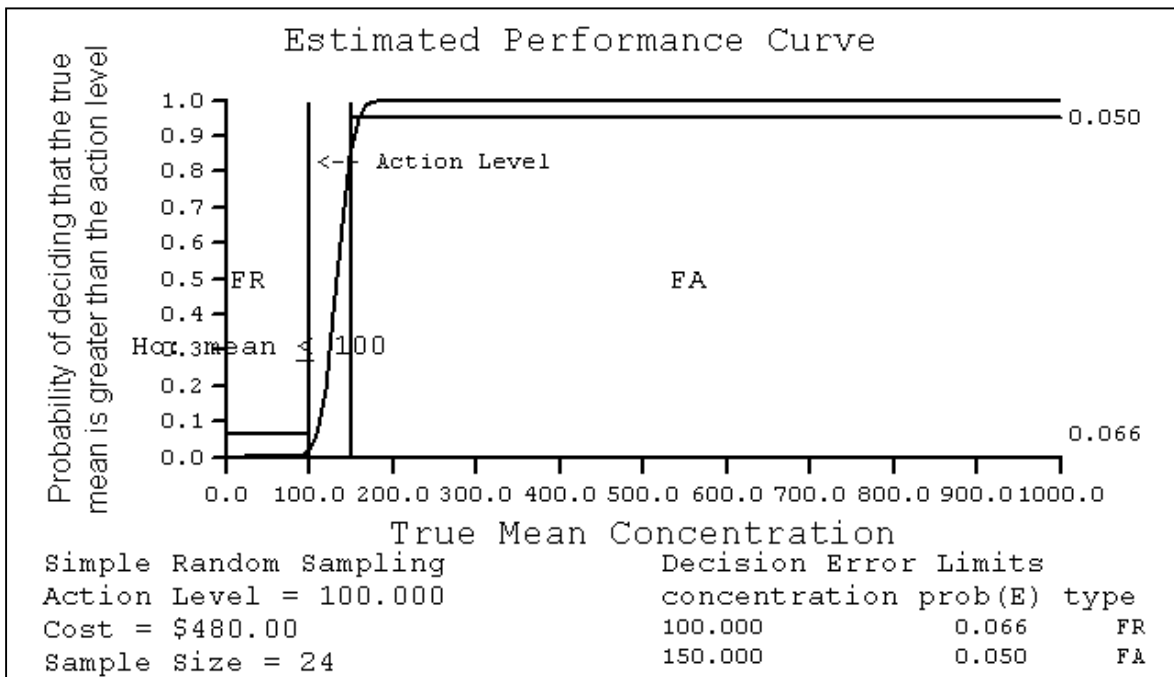
Note: The analysis is based on a coefficient of variation equal to 75 percent.

Figure 19
Routine to Concern Level Near-field River Section 2
(baseline to >60 mg/L with discrete samples every 3 hours for 6 hours)
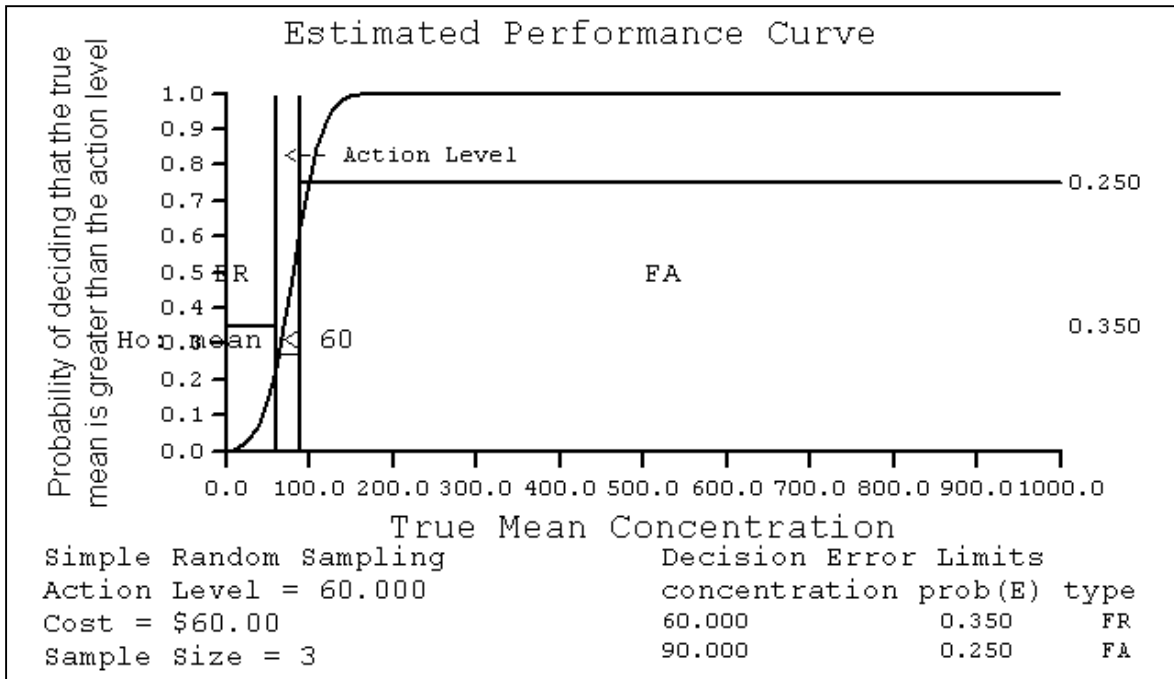
Figure 20
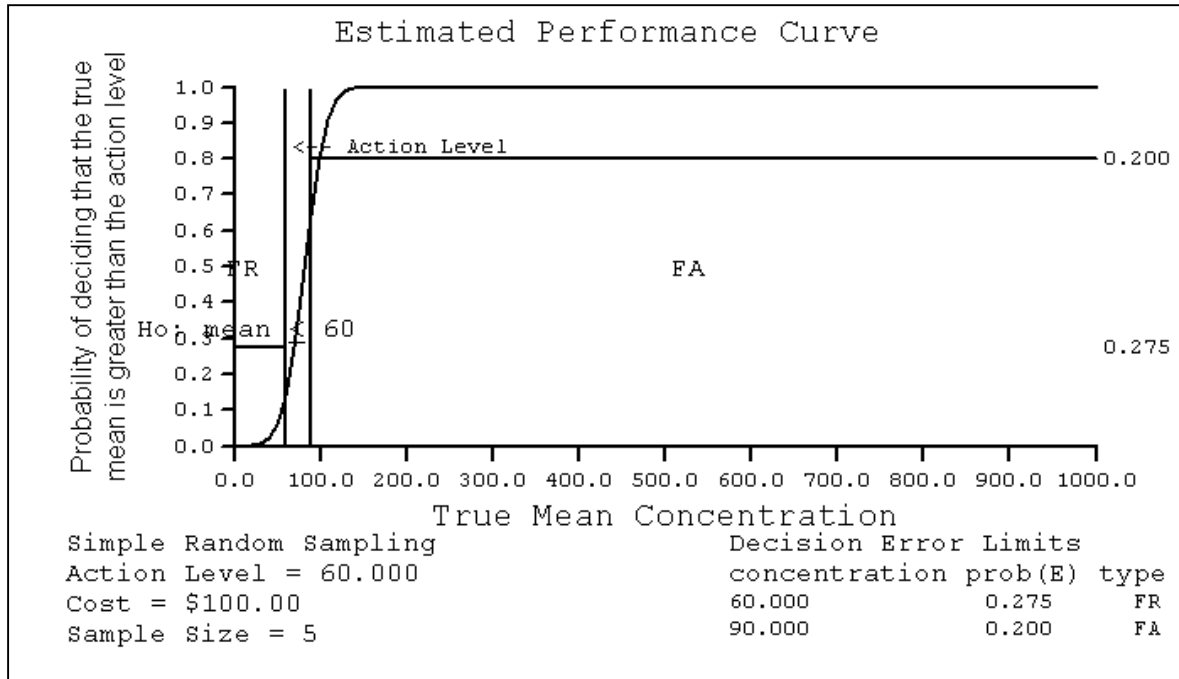**Routine to Concern Level Near-field River Section 2**
(baseline to >60 mg/L with continuous sampling every 15 min for 6 hrs)

Note: The analysis is based on a coefficient of variation equal to 75 percent.

Figure 21
Concern to Control Level Near-field River Sections 1 and 3
(baseline to >100 mg/L with discrete samples every 3 hours for 15 hours)

Figure 22
Concern to Control Level Near-field River Sections 1 and 3
(baseline to >100 mg/L with continuous sampling every 15 min for 15 hrs)

Note: The analysis is based on a coefficient of variation equal to 75 percent.

Figure 23
Concern to Control Level Near-field River Section 2
(baseline to >60 mg/L with discrete samples every 3 hours for 15 hours)

Figure 24
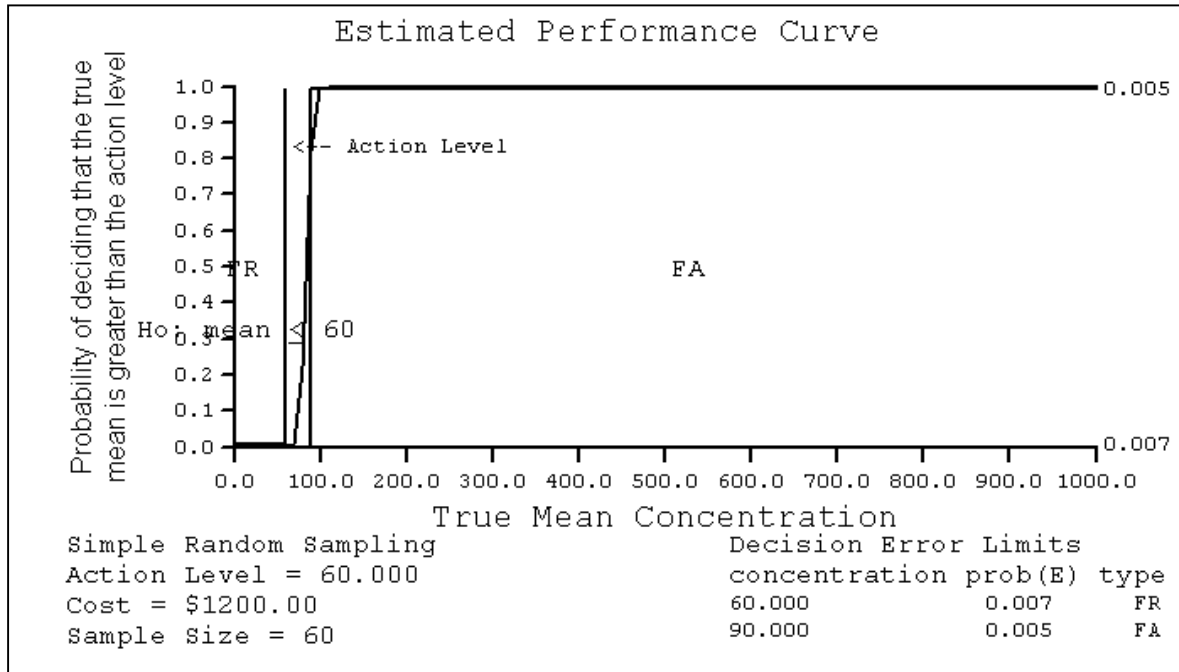**Concern to Control Level Near-field River Section 2**
(baseline to >60 mg/L with continuous sampling every 15 min for 15 hrs)

Note: The analysis is based on a coefficient of variation equal to 75 percent.

Figure 25
Routine to Evaluation Level
(Near-field baseline to >700 mg/L with discrete samples every 3 hrs for 3 hrs)
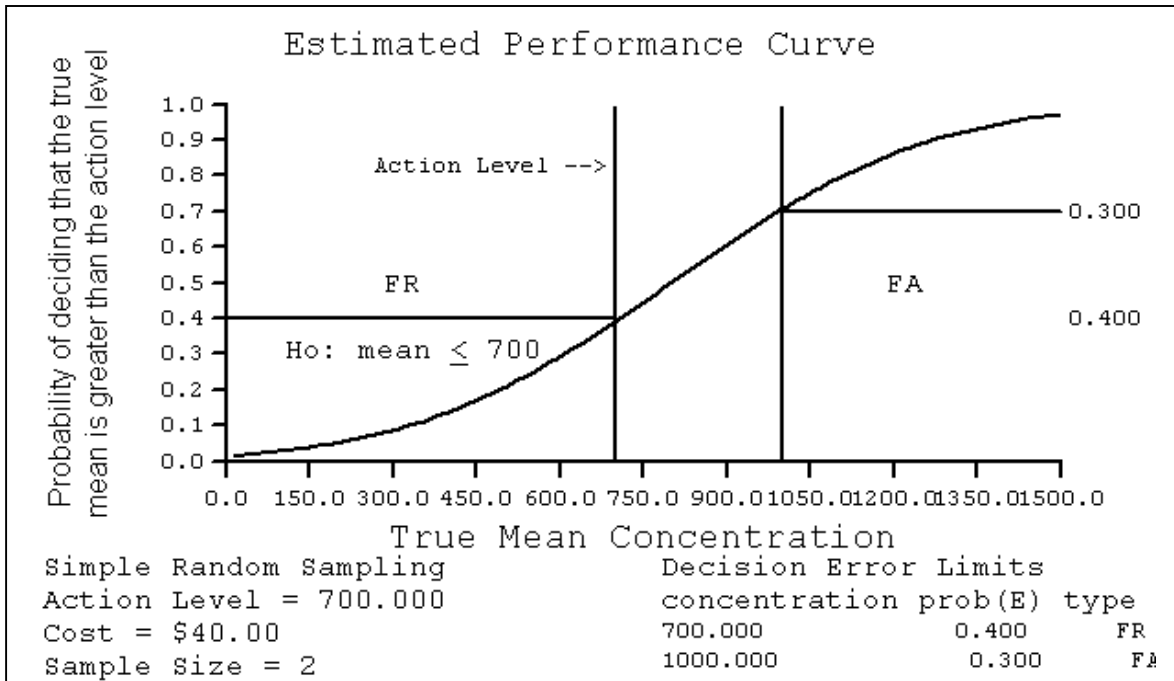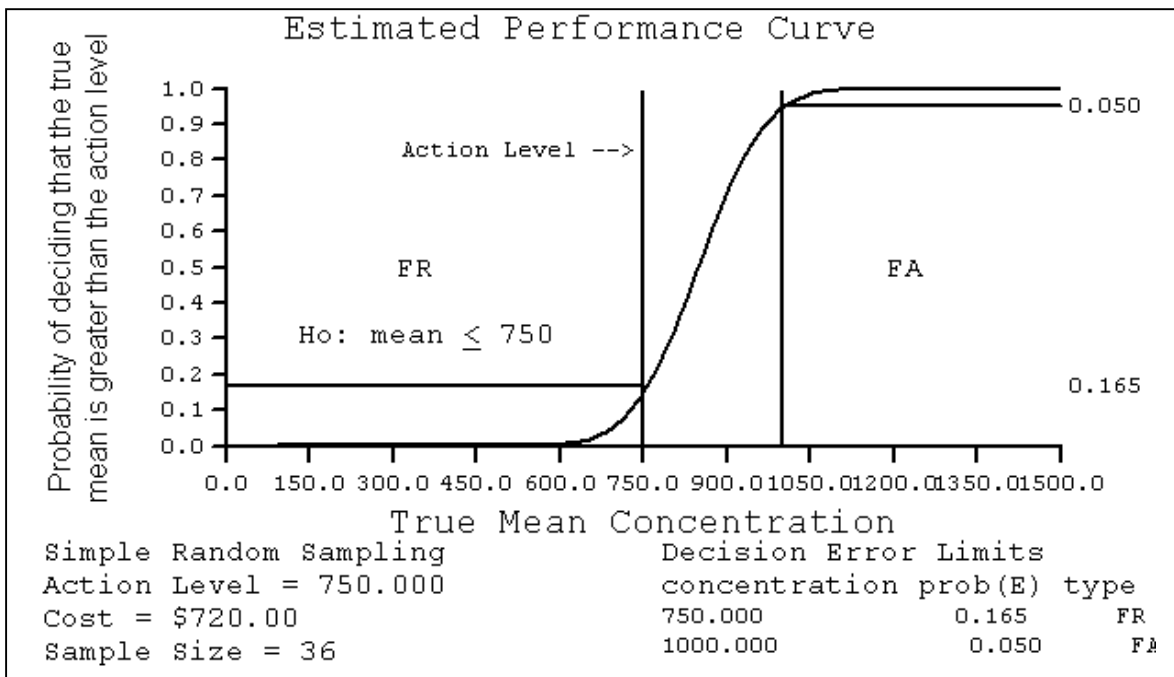

Figure 26
Routine to Evaluation Level
(Near-field baseline to >700 mg/L with continuous sampling every 15 min for 3 hrs)

Note: The analysis is based on a coefficient of variation equal to 75 percent.