



Standard Operating Procedures for Large Rivers Water Quality Monitoring

Great Lakes Network (Version 1.1)

Natural Resource Report NPS/GLKN/NRR—2016/1262.1



ON THE COVER

Preparing to lower a multi-probe sonde into the St. Croix River.
Photograph courtesy of the National Park Service/R. Damstra.

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Standard Operating Procedure #1: Pre-Season Preparation

Version 1.1

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Revision History Log

The following table lists all edits and amendments to this document since the original publication date. Information entered in the log must be complete and concise. Users of this standard operating procedure will promptly notify the project manager and/or the Great Lakes Network (GLKN) data manager about recommended and required changes. The project manager must review and incorporate all changes, complete the revision history log, and change the date and version number on the title page and in the header of the document file. For complete instructions, please refer to Revising the Protocol, SOP #13.

Previous Version Number*	Revision Date	Author (with title and affiliation)	Location in Document and Concise Description of Revision	Reason for Change	New Version Number
1.0	1/26/2016	David VanderMeulen, Rick Damstra. GLKN	updated the NPS contact author and authorship, edits to Table 2, updated the web link for obtaining a Research and Collecting Permit, added references to GLKN and park Safety Plans, fixed a handful of grammatical errors throughout	To clarify and update SOP based on changes that have occurred since Version 1.0 was published in 2008	1.1

*Add rows as needed for each change or set of changes tied to an updated version number

Acknowledgements

We relied upon documents by O’Ney (2005) and the U.S. Geological Survey (USGS 2005) for guidance in writing this standard operating procedure.

1.0 Introduction

Prior to the field season, many preparations must be completed to ensure sampling can be undertaken according to schedule. The field season for sampling large rivers of the Great Lakes Inventory and Monitoring Network (Network) parks is generally from March/April through November. All details for the season need to be planned well in advance. Field preparations should begin in January to allow enough time for ordering new supplies and equipment, if necessary. Table 1-1 provides a checklist and general guidance for activities conducted prior to sampling. Many of these activities are discussed in more detail in other SOPs.

Table 1-1. Checklist of activities to be conducted prior to sampling large rivers.

✓	Activity	Description
	Prepare calendar of planned field trips	Includes sampling dates, locations, personnel
	Review sampling methods to determine if revision is needed	Check web reference to see if method has been updated from version currently used
	Review checklists of equipment and supplies required; prepare list of items to be ordered	Check each Standard Operating Procedure for detailed equipment lists; check expiration dates of reagents, standards, and other chemicals
	Order supplies	Includes calibration standards, pre-cleaned bottles for sample collection (if not provided by the contract analytical laboratory) , sample preservation solutions
	Clean and test equipment; charge or replace batteries as needed	Includes multiparameter sonde, integrated sampling tube, Van Dorn, back-up instruments, ropes, thermometers, camera, GPS unit, cell phone or radio
	Ensure the latest versions of software are being used	Includes multiparameter sonde, data-logger, computer
	Prepare equipment blanks	Must be done annually
	Obtain permission for site access, if necessary	--
	Confirm current park research and collection permits	--
	Check field vehicle and boat for safety equipment and supplies	Includes material safety data sheets, flares, flashlight, gloves, extra sampling bottles, etc.
	Schedule boat training, if necessary	--
	Update field folder	Include maps, site information, field forms, sampling procedures
	Review sample collection, processing, and documentation information	Includes methods, lab codes, bottle type, and sample collection and processing procedures; sample shipment; quality control samples. (SOPs #6 and #7)
	Make travel reservations and arrangements as needed	--
	Provide supervisor with schedule	--

1.1 Read the Entire Protocol

Periodically read through the entire protocol, including all standard operating procedures (SOPs). Be sure to understand the purpose for which the various types of data will be collected and review the SOPs for the types of measurements and samples needed. Be alert for portions of the protocol or SOPs that may be in need of revision, and bring these sections to the attention of the appropriate supervisor.

1.2 Prepare Calendar of Planned Field Sampling

Well in advance of the field season, prepare a calendar of sampling dates for the entire season. Allow for the possibility of bad weather days, when sampling may have to be postponed. Include the location of sampling, dates, parameters to be measured, personnel, and any additional relevant notes (Table 1-2).

Table 1-2. Example of calendar of planned field sampling for large rivers park.

Dates	WQ Variables	Personnel	Notes
April 4-6	core suite, chl-a, TP, TN, alk, NO ₂ +NO ₃ -N, NH ₄ -N, TSS, Alk, Cl, SO ₄ , DOC, Cations, SiO ₂	Damstra, park biotech	Quarterly sampling of ions, etc.
May 2-4	core suite, chl-a, TP, TN, alk, NO ₂ +NO ₃ -N, NH ₄ -N, TSS	Damstra, park biotech	--
May 31-Jun 2	core suite, chl-a, TP, TN, alk, NO ₂ +NO ₃ -N, NH ₄ -N, TSS	Damstra, park biotech	--
July 11-13	core suite, chl-a, TP, TN, alk, NO ₂ +NO ₃ -N, NH ₄ -N, Bottom TP, TSS, Alk, Cl, SO ₄ , DOC, Cations, SiO ₂	Damstra, park biotech	Quarterly sampling of ions, etc.; collected bottom water sample if stratified
August 8-10	core suite, chl-a, TP, TN, alk, NO ₂ +NO ₃ -N, NH ₄ -N, TSS, cations, anions, DOC, SiO ₂	Damstra, park biotech	--
Sept 12-14	core suite, chl-a, TP, TN, alk, NO ₂ +NO ₃ -N, NH ₄ -N, TSS	Damstra, park biotech	--
October 3-5	core suite, chl-a, TP, TN, alk, NO ₂ +NO ₃ -N, NH ₄ -N, TSS, Alk, Cl, SO ₄ , DOC, Cations, SiO ₂	Damstra, park biotech	Quarterly sampling of ion, etc.
November 7-9	core suite, chl-a, TP, TN, alk, NO ₂ +NO ₃ -N, NH ₄ -N, TSS	Damstra, park biotech	--

1.3 Review Checklists of Equipment and Supplies

Checklists help ensure that equipment and supplies will be ordered on time, data collection activities will be completed appropriately, and data quality objectives will be met. Review the detailed equipment lists that are included with each standard operating procedure (Table 1-3). Pay attention to

expiration dates on reagents, calibration standards, and all other chemicals. Prepare a list of equipment and supplies that must be ordered and present it to the project manager.

Table 1-3. Checklists of equipment and supplies for monitoring water quality of large rivers.

Checklist	Location
Safety equipment checklist	SOP #2
Decontamination equipment and supplies	SOP #5
Field supplies and equipment checklist	SOP #6
Laboratory equipment and supplies	SOP #7

1.4 Confirm/Apply/Renew Research and Collecting Permits

For sampling stations located within park boundaries, a National Park Service Research and Collecting Permit must be obtained before any work can be done. To renew or apply for the permit, go to the following website: <https://irma.nps.gov/rprs/>

Follow the directions for renewing an existing permit, or if your project is not already in the system, then follow the directions to apply for a new permit. Work with the research coordinator at each park.

1.5 Review Sample Collection, Processing, and Documentation

Conduct a thorough review of “Field Measurements and Water Sample Collection” (SOP #6) and “Processing Water Samples and Analytical Laboratory Requirements” (SOP #7).

Check with the project leader to determine current contract analytical laboratory information. Contact the laboratory to verify lab codes and procedures. Obtain copies of Analytical Services Request (ASR) and Chain of Custody (COC) Forms.

1.6 Update Field and Office Binders

Field binders should contain reference information specific to each sampling station, including maps, photos, previous data, field forms, and summaries of sampling and quality assurance/quality control (QA/QC) procedures. The office binder should contain reference information relevant to general field sampling procedures, including QA/QC reminders, copies of all SOPs relating to safety, decontamination, sample collection and processing, copies of equipment instructions and troubleshooting, calibration logs, Safety Data Sheets (SDSs), and ASR and COC forms. The field binder should be taken along on each sampling trip.

Each year, prior to the sampling season, the field binder for each monitoring site should be reviewed and the following information updated as needed:

- *Location of river level gage (if one is present).*
- *Location of sample-collection sites.* Review field notes for any indication that the location for sample collection may need revision. Update protocol if necessary.
- *Name of landowner, tenant, or other responsible party.* If the sampling station is located on private land, ownership may change. Verify.
- *Current copy of research and collection permit (if site located within NPS boundaries).* Check dates on permit. Renew/apply as described above.
- *Site access instructions (for example, call owner or site operator before arrival at site, obtain key to unlock security gate).* Confirm contact person, procedure, and phone numbers.
- *Photographs to document site conditions.* Take new digital photograph annually.
- *Maps to site (state and local).* Review map for accuracy; update if necessary.
- *Review previously collected chemical, physical, and biological data.* A summary of previously-collected data or copies of previous field data sheets should be in the field folder so that water column profiles and Secchi disc/transparency tube data can be compared to previous surveys. Familiarity with previously collected data is a critical quality assurance (QA) element for early detection of possible instrument malfunction.
- *Summaries of field procedures, QA/QC procedures, instrument stabilization criteria, etc.* Prepare brief outlines reminding field personnel of routine procedures, including all changes from the previous year.

Each year, prior to the sampling season, the office binder should be reviewed and the following information updated as needed:

- *Safety information (SOP #2).* Verify/update “Medical Information Form for Field Personnel” and “Emergency Contact Form”.
- *Sampling schedule and instructions.* Ensure the following information from SOP #6 (Field Measurements and Water Sample Collection) is included in the office binder: laboratory analyses to be requested and associated codes, when to collect samples, bottle types needed for each analytical schedule, preservation requirements, quality control sample requirements, and shipping instructions. Verify information is correct and report and reconcile any discrepancies to supervisor.
- *Decontamination procedures (SOP #5).* Ensure that the most recent information on presence of exotic species and need for decontamination of equipment has been incorporated into the Hazard Analysis and Critical Control Plan (HACCP) for each park. Update office binder with the latest information. If the order of lakes to be sampled must change due to new infestations, be sure sampling schedule is updated.

- *Analytical service request forms, data collection field forms, chain of custody forms, sample bottle labels.* Prepare as much of the field forms as possible in advance. For each station, complete the header information, including the project ID, station ID and station name and other required metadata for NPSTORET. Place enough blank field forms in office folder to last entire field season.
- *Ensure that copies of the current field procedures are included in the office folder.* Include calibration and maintenance procedures specific to the instruments to be used.

1.7 Clean and Test Equipment

Clean and test all sampling equipment, including multiparameter sonde, temperature sensors, Van Dorn and integrated tube samplers, camera, GPS units, and any back-up meters. Check calibration of metered ropes to ensure accurate depth measures. Start each new field season with fresh batteries and replace spares in field tool kit. Ensure the latest versions of software are loaded onto the multi-probe, data-logger, and computer.

1.8 Vehicle, Boat, and Safety Gear

Check maintenance schedule of field vehicle and arrange maintenance, if needed. Check boats and vehicle for safety equipment such as MSDS sheets, flares, spare tire, triangles, cones, first aid kit. Prepare a list of supplies needed and present list to supervisor. If using a trailer, ensure that tail-lights are in working order. Check that field tool-kit is complete and replace tools, as needed.

1.9 Training and Safety

Keep current with training and the laboratory requirements associated with your data collection activities. New technicians will need basic skills training, including classroom instruction, hands-on training, and pilot-testing of equipment. Continuing field staff should attend refresher training and review Network and park safety plans annually. If boat training is needed, be sure to schedule the training with the park early in the season. Initial and periodic refresher courses in basic first aid and CPR are required.

1.10 Field Reconnaissance

Make field reconnaissance trips, if possible. Visit the sampling sites to be sure that conditions have not changed from the previous year. Note conditions that could affect sampling operations, such as the seasonal high or low water levels, or site access peculiarities, and if necessary investigate alternative access points to launch a boat, for example.

1.11 Travel Arrangements

Make travel arrangements. Because hotel reservations may be difficult to impossible to obtain at certain times of year, it is important to review the sampling schedule and plan ahead. If necessary, help coordinate travel for field staff.

1.12 Communicate with Supervisor

Ensure the project manager is informed of supply needs, problems with instruments, changes in sampling schedule, changes in sampling site conditions, and other needs that may have an impact on the project budget, data collection, schedule, or sampling design.

1.13 Equipment Blanks

Equipment blanks should be conducted annually at each park where sampling is scheduled, at least four weeks prior to using the equipment in the field to ensure adequate time for analysis and review of results. Equipment blanks should be collected in a designated clean area of the sample processing laboratory. The blank consists of deionized water that is passed sequentially through each component of the sample collection and processing equipment. Equipment blanks should also be conducted when a new cleaning procedure is instituted and when new equipment will be used for the first time.

Procedure

- Fill the integrated sampling tube or Van Dorn with deionized water (DIW).
- Dispense water from integrating tube or Van Dorn into cubitainer or carboy.
- From cubitainer or carboy, dispense water directly into appropriate analyte bottles, or filter first, according to instructions for each analyte (SOP #7).
- Preserve samples according to instructions for each analyte in SOP #7.

If the equipment-blank data indicate that the equipment does not introduce contaminants that will bias study results, sampling can proceed. If the equipment-blank data indicate unacceptable concentrations of analytes of interest, the cause must be identified and the equipment or cleaning procedures must be changed or modified before sampling can proceed.

1.14 Literature Cited

- O'Ney, S. E. 2005. Standard operating procedure #2: Pre-season activities, Version 1.0. *In* Regulatory water quality monitoring protocol, Version 1.0, Appendix E. National Park Service, Greater Yellowstone Network, Bozeman, MT.
- USGS (U.S. Geological Survey). 2005. National field manual for the collection of water-quality data: U.S. Geological Survey Techniques for Water-Resources Investigations, book 9, chaps. A1-A9.

Standard Operating Procedure #2: Training and Safety

Version 1.1

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1.0	1/27/2016	David VanderMeulen, Rick Damstra, GLKN	Updated the NPS contact author and authorship, added a number of sections taken directly from the Great Lakes Network Health and Safety Plan, included use of a stadia rod and hand level to measure water level, fixed a handful of grammatical errors throughout	To clarify and update SOP based on changes that have occurred since Version 1.0 was published in 2008	1.1

*Add rows as needed for each change or set of changes tied to an updated version number

Acknowledgements

The safety related portions of this standard operating procedure were adapted from the Great Lakes Network Health and Safety plan (Route et al. 2013) and the Greater Yellowstone Network's SOP on safety (O'Ney 2005), with heavy reliance on the U.S. Geological Survey's (USGS) manual for collecting water quality data (Lane and Fay 1997).

2.0 Introduction

Prior to collecting data and water samples in the field, the field crew members must be trained in the techniques and procedures that will be used throughout the season. Familiarity with the protocol and standard operating procedures (SOPs), as well as with boats, equipment, and basic safety standards, are critical to the success of the water quality monitoring program. This SOP includes procedures for training in the specific water quality related skills and knowledge necessary for collecting good data and understanding those data. The crew must be also trained in specific safety procedures to ensure their safety and that of others.

The project manager will conduct or arrange for all training needed prior to the field season.

2.1 Pre-Season Classroom Training

Periodically read through the entire protocol, including all standard operating procedures (SOPs). Be sure to understand the purpose for which the various types of data will be collected and review the SOPs for the types of measurements and samples needed. Be alert for portions of the protocol or SOPs that may be in need of revision, and bring these sections to the attention of the appropriate supervisor.

It is desirable to begin training well before the field season begins to allow adequate time for thorough understanding of field and laboratory procedures and to obtain certification in boat use. Field crew leaders must undergo all of the following training. Training in boat use is highly recommended for other field crew members, and training in the remaining areas is desirable.

2.1.1 Limnological Concepts

An understanding of basic limnological concepts is useful for recognizing bad or illogical data in the field. For example, a dissolved oxygen reading of 30 mg/L may indicate a malfunction of the multiparameter sensor, or something as simple as fouling of the probe. Recognition of the problem at the time it occurs allows for immediate adjustment in the field, for example cleaning the membrane on the probe or changing the batteries, so that good data can still be collected.

The field crew leaders, and if possible, other crew members, will study select modules of the Water on the Web (2004) curricula (<http://waterontheweb.org/curricula>). The modules will be selected by the project manager and will include those on lake surveys (e.g., field profiles, sample collection, laboratory methods). Individual study will be followed with group discussion and/or individual discussion with the project manager.

2.1.2 Understanding the Protocol and Standard Operating Procedures

Reading and understanding the entire protocol and all SOPs are crucial prior to initiating field work. The project manager will allow adequate time for all field crew members to complete this step to ensure success of the project. Field and laboratory related SOPs will also be covered as part of the hands-on training, described below.

2.1.3 First Aid and Cardiopulmonary Resuscitation (CPR)

Training in the basic medic/first aid and Heartsaver AED, which includes CPR and use of an automated external defibrillator (AED), is required for all crew members (Route et al. 2013) and will be paid for by the Network. Acceptable training should be through the American Red Cross or American Heart Association. Certification is valid for two years. Training and certification should be acquired prior to the field season.

2.2 Hands-On Training

In addition to the classroom training described above, a variety of hands-on training and practice prior to the first sampling period will help ensure high quality data collection. Familiarity with the use and maintenance of equipment, procedures for collecting and processing water samples, techniques for cleaning field and laboratory equipment, and safe use of watercraft are essential to the success of the water quality monitoring project. Field crew leaders are required to complete all of the following training; other field crew members should also complete the training, if possible, although it is not required.

2.2.1 Use of a Multisensor Water Quality Instrument

Each type of multisensor instrument comes with specific instructions on use and care. Guidelines for calibration and use of multisensor instruments are included in SOP #6, however it is important to use the instruction manual specific to each instrument. Check the manufacturers' websites for updates or changes to instructions.

Training in the use of a multisensor instrument will include the following:

- calibration procedures and acceptance criteria
- keeping a calibration log
- maintenance, including replacing fluids and membranes
- creating files on the data logger
- storing data on the data logger
- downloading data to a laptop computer
- use of the instrument in a field setting, including depth profiles and equilibration
- troubleshooting

Crew members will have the opportunity to practice the skills learned prior to the actual sampling until they are comfortable with the use of the instrument.

2.2.2 Use of a Global Positioning System (GPS)

Location information must be gathered via GPS for each site during each visit. Training in the use of a GPS will include navigation to a known location, acquiring location information, storing data, and downloading data. Details on the use of a GPS can be found in SOP #3.

2.2.3 Field Methods

In addition to collecting data with the multisensor instrument, a field data sheet must be completed, a water sample must be collected, and where possible the water level must be measured. Prior to the field season, the field crew will receive training in and have the opportunity to practice the following:

- completing the field data sheet
- using an integrated sampling tube
- using a Van Dorn sampler
- measuring water level by reading a measuring tape with a plumb bob or a staff
- decontaminating equipment between lakes to avoid transfer of species from one area to another, and determining when decontamination is necessary
- following QA/QC procedures

2.2.4 Processing Water Samples

Whether the water samples are processed for further analytical laboratory analysis in the field or back at the office or lab, strict procedures must be adhered to (see SOP #7). Crew members will be trained in, and will have the opportunity to practice the following techniques:

- handling water samples so as to avoid contamination
- rinsing and filling bottles from the analytical laboratory
- preserving samples for various chemical analyses
- filtering samples for various analyses
- packing samples for shipment to analytical laboratory
- filling out chain of custody forms
- following QA/QC procedures

2.2.5 Cleaning Field and Laboratory Equipment

Field and laboratory equipment must be cleaned between lakes or samples to avoid contamination with water from the previous sampling site. Training in proper techniques will include:

- cleaning equipment with phosphorus-free detergent
- setting up an acid bath
- cleaning equipment with an acid bath
- rinsing with distilled or de-ionized water

2.2.6 Boat Training

Prior to operating a NPS boat or canoe, training and certification are required (Route et al. 2013). The crew leader must receive the training and obtain certification, and it is highly desirable for the remaining crew members to do so as well. The project manager will arrange for training prior to the

field season through one of the several GLKN parks that offer it. Arrangements should be made well in advance of the field season.

2.3 Safety Procedures

Safety of personnel should always be the first concern in conducting a monitoring program and in the selection of sampling sites. General safety considerations for work in the office, field, and laboratory, as well as a discussion of other issues such as prevention of accidents and injuries and communication procedures are discussed in the Great Lakes Network Health and Safety Plan (Route et al. 2013) and GLKN's Laboratory Safety Plan.

Numerous safety issues and concerns are associated with implementing a water quality monitoring program that includes extensive field work and sampling. Field personnel routinely come into direct and indirect contact with waterborne pathogens, chemicals, and potentially hazardous plants and animals. Field work requires an awareness of potential hazards (i.e., situational awareness) and knowledge of basic safety procedures. Advanced planning can reduce or eliminate many safety hazards.

2.3.1 USGS Field Manual

This SOP is meant to be used in conjunction with Chapter A9 of the USGS National Field Manual (Lane and Fay 1997), which contains more complete information about potential hazards that water quality monitoring field personnel may encounter during field work and the procedures that, when implemented properly, will help ensure the safety and health of field crew members. A copy of this manual is provided to the field crew and may be downloaded from <http://water.usgs.gov/owq/FieldManual/Chap9/content.html>. Topics addressed in the USGS document include:

- general references for federal policies and Department of Interior (DOI) safety guidelines
- safety policies you are required to know and follow under the Occupational Safety and Health Act (OSHA), Environmental Protection Agency (EPA), and Department of Transportation (DOT);
- understanding and implementing a job hazard analysis (JHA);
- requirements related to use of personal protective equipment (PPE) on the job
- safety training and certification requirements; safety issues associated with transportation and operation of vehicles (road vehicles and trailers, watercraft, aircraft etc.) used to reach sampling sites
- surface water activities (e.g., wading, working from bridges, boats and cableways, etc.)
- working around machinery, pumps, and other equipment
- proper use, handling, transport, storage, and disposal of chemicals
- handling of contaminated water and limiting exposure to yourself and others

- environmental conditions caused by extremes in temperature; sun exposure; threats posed by storms, floods, fire, snow, ice, and various animals and plants

In addition to consulting the USGS manual, the field crew should contact individual park's safety officers or resource managers for information on park safety plans, radio safety procedures, and local problems and issues, such as dangerous or nuisance animals, insect-and tick-borne diseases (e.g., Lyme disease, encephalitis, West Nile disease), and other issues specific to each park.

2.3.2 Basic Fieldwork Safety Preparation

2.3.2.1 Crew Size

Basic preparations for fieldwork should become routine before every sampling activity. The minimum recommended crew size for field projects is two people. Working alone in the field may be allowed if authorized in advance by your supervisor. If it is absolutely necessary for a person to work in the field alone, this person must:

- Have verbal authorization from your supervisor
- Carry two forms of reliable communication (e.g. radio and cell phone)
- Communicate periodically with co-workers or radio dispatch about planned travel routes, and times of departure and return to key locations.

2.3.2.2 Safety Gear and Information

Carry basic safety equipment, including first aid kit, flashlight, boots, rain gear, antibacterial soap or hand cleaner, matches or lighter, etc. Be aware of changing weather conditions and the potential for storms. Be aware of potential hazards at a monitoring site. Carry general safety information in each vehicle or boat, including:

- safety data sheets (formerly referred to as material safety data sheets) for preservatives
- basic first aid protocols
- emergency phone numbers
- locations of emergency facilities (hospitals, police and fire departments, U.S. Coast Guard)
- maps of the park, surrounding area, and nearest city

Job hazard analyses (JHAs), prepared by the aquatic ecologist, will be discussed with field personnel prior to the field season. The JHAs will cover such topics as hiking and portaging, boating and sampling from a boat/canoe, lab safety, driving vehicles, and stinging insects and poisonous plants.

The next two topics, Trip Plans and Methods of Communication, cover many aspects of fieldwork preparation, and are nearly verbatim from the Great Lakes Network Health and Safety Plan (Route et al. 2013).

2.3.2.3 Trip Plans

For safety purposes, it is critical that certain park staff be aware of fieldwork being conducted in their park by the Network. Network employees are required to set up a chain of communication and provide the park with essential information regarding Network activities—herein referred to as a “Trip Plan.” For front-country fieldwork this can be as simple as an informal email to the host park’s primary contact and cc’d to your supervisor. All trip plans must include who will be involved in the fieldwork, when the fieldwork will take place, what mode of transportation you have, where the work will be performed, and how you can be communicated with. For example an informal email:

To: Park Contact

cc: Joe Supervisor.

I and Jane Doe will be in your park tomorrow, 6/15/2013, to conduct water quality monitoring at Pristine Lake from approximately 10:00 AM until 2:00 PM. We will park our NPS vehicle at the boat ramp for Big Lake, boat to the dock at Grassy Campground, and portage a canoe to Pristine Lake using Rocky Portage Trail. I will have a cell phone (###-###-####) and a park radio, but the radio will be off unless I have an emergency.

Such an email is not only a good way of providing essential safety information, it is good policy for keeping the park informed of our activities.

For all backcountry trips, or for fieldwork spread over several days, a more formal trip plan should be filed. The Network program manager is responsible for ensuring the trip plan meets the host park’s need for detail and flow of communication. Refer to Appendix H of the Great Lakes Network Health and Safety Plan (Route et al. 2013) or the Network SharePoint site under the Safety folder for an example template of a trip plan.

All trip plans should consider the following guidelines and standards:

- A trip leader must be identified when more than one person is involved.
- The trip leader must file a trip plan with the host park and his/her supervisor or delegate.
- The supervisor is responsible for the flow of communication internally within the Network.
- A primary contact will be established with the host park through the park’s chief ranger or superintendent. This is the responsibility of the project supervisor, but may be coordinated more broadly by the Network program manager. The chief ranger or superintendent may delegate a primary contact.
- The park’s primary contact is responsible for establishing the park’s internal communications, ensuring Network trip plans are available to park staff, and if necessary, initiating an emergency response according to park policy. In parks where there is a dispatcher, it is the responsibility of the park’s primary contact to see that all dispatchers are aware of Network activities.

- The park resource manager/biologist will be informed of Network activities and is often delegated as the primary contact.
- For backcountry trips, the trip leader will post the trip plan on the Network SharePoint site under the Safety folder (<http://imnetsharepoint/glkn/safety/default.aspx>). Depending on the complexity and inherent dangers of the fieldwork, supplemental information like GPS coordinates for sites or maps of survey areas will also be posted to the SharePoint.
- Check-ins may be required by the park. If required, the timing and frequency should be documented in the trip plan.
- One trip plan can cover an entire field season, but the plan should clearly outline the inclusive dates of discrete field efforts.
- If plans change to such a degree that someone would be unable to easily locate the crew, the trip leader should inform the park as soon as practicable.

2.3.2.4 Methods of Communication

Field crews must have at least two methods of communication and should determine the most effective methods in consultation with their supervisor and the park prior to the trip. The methods will vary by park and situation, but the most common will be a portable two-way park radio (for parks using radio systems) and either a cell phone or a satellite phone. For urban parks, this can be two cell phones (best if they have different carriers to increase the chance of reception), or a cell phone and a satellite phone. Park staff should be consulted regarding ‘dead zones’ in radio and cell coverage. All GLKN vessels on the Great Lakes will have a park radio and a marine-band radio mounted in the cabin.

For remote situations on land, crew leaders will also carry a Spot Locator or equivalent device. In the event a radio is not working properly it can be used to send an email to select individuals and/or request immediate emergency response. For boating on the Great Lakes, all GLKN vessels over 18 feet will be equipped with an emergency position-indicating radio beacon (EPIRB). The EPIRB is preprogrammed to send an emergency message to the U.S. Coast Guard.

SOPs for contacts, using portable radios, Spot locator devices, and satellite phones are available within the Great Lakes Network Health and Safety Plan (Route et al. 2013). Supervisors are required to provide hands-on training with crew members on use of devices that will be used during their field work.

2.3.3 Emergency Medical Forms and Information

In a true emergency, get the patient to medical care immediately and worry about forms, payments, and other procedures after the situation has been stabilized. For all non-emergency medical treatments, you must have your supervisor authorize medical treatment and complete a CA-16 form (Authorization for Exam/Treatment) and submit this form to the medical provider. If your supervisor is unavailable and you cannot wait for the treatment, call the Network office or the park and someone should be able to authorize treatment.

Accidents and/or injuries that occur on the job must be reported on either a CA-1 (Federal Employee's Notice of Traumatic Injury and Claim for Continuation of Pay/Compensation) or CA-2 (Federal Employee's Notice of Occupational Disease and Claim for Compensation) filled out on the Safety Management Information System (SMIS) website: <https://www.smis.doi.gov>. The injured employee fills out the form online and the supervisor is notified by email that his/her portion of the claim is ready to be filled out. Once completed by the supervisor, the claim is forwarded to the Midwest Region claims coordinator who reviews and forwards the claim to the Office of Worker's Compensation Program (OWCP).

The following paper forms can aid in gathering information for reporting an accident and/or injury that occurred on the job and can be obtained from the U.S. Office of Personnel Management (OPM) website (<http://www.opm.gov/>).

- You may need to provide OWCP1500 "Health Insurance Claim Form" to your provider.
- If you require time off work you must complete form CA-17. Your supervisor must fill out the first portion describing your work duties. The treating physician should then complete the second portion of the CA-17, stating what tasks you cannot perform.
- For reference see CA-10 "What a Federal Employee Should Do When Injured At Work."

Employees have a reasonable expectation of privacy; however, the NPS is responsible for assisting employees in the event of a medical emergency. To do so, we must be aware of allergies, prescription medications, or other medical conditions that would help emergency medical technicians and physicians perform their duties. All Network employees have the option to complete a Medical Information Form (Appendix 1) that informs their supervisor of medical conditions that could affect their health and well-being. Employees who prefer not to share this information must still sign and submit a Medical Information Form to their supervisor, indicating their preference. The original copy will be placed in a sealed blue envelope by the employee and kept in a locked drawer in the Network administrative office. For employees who are in the field, a copy of this form will be kept by the supervisor or crew leader in a sealed blue envelope. These sealed envelopes will be opened only in an emergency. They will be shredded by the employee at the end of employment.

2.3.4 Safety Equipment Checklists

Checklists are helpful for ensuring that personnel have the appropriate safety equipment available during field trips. Items shown in Tables 2-1 through 4 are the basic necessities for field work. Field crew members should consider their specific needs and should customize the checklists as necessary. The field crew and project manager will discuss the checklists and determine which items are necessary.

Personal Protective Equipment (PPE) (Table 2-2) must be selected based on the hazards likely to be encountered. The Great Lakes Network is required to supply appropriate PPE, and field personnel are required to use it.

Table 2-1. General safety equipment for field work.

√	Basic Safety Equipment Checklist
	Waders, hip boots, rubber knee boots
	Personal floatation device (PFD)
	First aid kit
	Fire extinguisher
	Flashlight and spare batteries
	Communication devices (Park radio, cellular or satellite phone, SPOT locator)
	Rain gear
	Hat, sun screen, and sunglasses
	Drinking water or sports drinks
	Safety cones, orange safety vest (working on bridges)
	Tool box with basic tools
	Antibacterial soap or hand cleaner
	Spill kits (for preservatives)
	Material safety data sheets (MSDS) for preservatives
	Hand-held eye wash
	Protective safety glasses or goggles
	Container to carry preservatives
	List of emergency phone numbers and office contacts

Table 2-2. Personal protective equipment for laboratory and field work.

√	Chemical and disease protection
	Aprons
	Eye/Face splash guards
	Gloves (vinyl and/or latex or nitrile)
	Protective suits
	Respirators (certification required for use)
√	Weather and UV protection
	Boots
	Fluids (e.g., water, sports drinks)
	Hat with a brim
	Insect repellent
	Rain gear
	Sunglasses
	Sunscreen
	Temperature-modifying clothing
	Work gloves
√	Flotation and reflective protection
	Orange flotation vests and jackets
	Safety harness
√	Protection for working around boat motors
	Hearing protection

Table 2-3. Checklists for vehicles and vehicular laboratories.

√	Chemical protection and storage
	Chemical spill kit
	Eye wash kit (replace old or expired wash solution)
	Safety Data Sheets (SDS)
	Chemical reagents (stored in appropriate area)
	Flammable solvents (stored in appropriate dedicated area)
	Pressurized gases (stored in appropriate area)
√	Communications and instructions
	Field folder (including maps, emergency phone numbers for medical facilities, office contacts, family contacts)
	Cellular phone/communication equipment (check that the service is operational for the area to be traveled)
√	First aid and protective equipment
	Complete change of clothes (stored in dry area)
	Fire extinguisher (safely secured)
	First aid kit and manual (check for missing or old, expired items and replace if necessary)
	Orange reflective vest
√	Miscellaneous equipment
	Bungee cords (to secure loose articles)
	District flood plan (most current version)
	Flagging
	Duct tape
	Knife or multi-tool
	Flares
	Flashlight (including fresh batteries)
	Flexible hose (to vent exhaust away from vehicle)
	Safety cones
	Tool kit
	U.S. Geological Survey TWRI Book 9 Chapter A9.

Table 2-4. Checklists for the use of watercraft.

✓	Instructions and navigation
	Field folder, with sampling plans
	Charts and maps
	Compass
	Depth finder
	Dead-man's switch
	Navigation lights
	Ring buoy with line
✓	Distress and external communication
	Radio (VHF, AM, FM, and WEATHER)
	Special lighting/flagging (if boat activities might pose a hazard to the public, such as tag line measurements)
	Visual distress signals (Coast Guard approved)
	Whistles or horns
	Type IV throwable rescue device
	Personal flotation devices for each passenger (Coast Guard approved)
	Anchor and lines (spare)
	Bucket for use as a bailer (sponge for use in canoes)
	Paddle (extra paddle for each canoe or rowboat)
	First aid kit (Coast Guard approved)
	Flashlights and batteries
	Fire extinguishers
	Spare parts (anchor, fuel, propeller, extra lines, cotter pin)
	Tool and repair kits
	Extra clothes (hat, foul-weather gear)
	Food and water
	Sunscreen
	Conversion factors and abbreviations

2.4 Literature Cited

- Lane, S. L., and R. G. Fay. 1997. Ch A9: Safety in field activities. *In* National field manual for the collection of water-quality data: U.S. Geological Survey techniques of water-resources investigations, Book 9. Available at <http://pubs.water.usgs.gov/twri9A9>.
- O’Ney, S. E. 2005. Standard operating procedure #3: Safety and health, Version 1.0. *In* Regulatory water quality monitoring protocol, Version 1.0, Appendix E. Bozeman (MT): National Park Service, Greater Yellowstone Network.
- Route, B., C. Chapin, T. Gostomski, and I. Messerly. 2013. Great Lakes Network health and safety plan, version 1.0. Unpublished document. National Park Service, Great Lakes Network Office, Ashland, Wisconsin.
- Water on the Web (WOW). 2004. Monitoring Minnesota lakes on the internet and training water science technicians for the future—A national online curriculum using advanced technologies and real-time data. University of Minnesota–Duluth, Duluth, Minnesota. (<http://waterontheweb.org/curricula>).

Appendix 2A. Emergency Contact and Medical Form

Employee Name _____

Emergency Contacts

#1 Name: _____ Relationship: _____
Phone: (home) _____ (work): _____

#2 Name: _____ Relationship: _____
Phone: (home) _____ (work): _____

Great Lakes Network Contacts

Network Office	715-682-0631 x224	Mississippi NRRRA	_____
Apostle Islands NL	_____	Pictured Rocks NL	_____
Grand Portage NM	_____	St. Croix NSR	_____
Indiana Dunes NL	_____	Sleeping Bear Dunes NL	_____
Isle Royale NP	_____	Voyageurs NP	_____

Local Emergency Contacts (or call 911)

Nearest Hospital:

Address: _____
Phone: _____

Other medical facility (24-hour care)

Address: _____
Phone: _____

Police: 911 or _____

Fire: 911 or _____

Utility: _____

Health Information Centers

Poison Control: 1-800-222-1222 _____

Centers for Disease Control: _____

Information Hotline: _____

Other: _____

Medical Information Form (*retain in office*)

Name

Date of birth

Local mailing address

Local phone number

Cell phone number

Emergency contact
information #1

Relationship:

Emergency contact
information #2

Relationship:

Allergies and medical
issues of concern

Medications currently
being taken

☐ Check here and provide additional information on the back or on attached sheets if there are other issues you feel emergency personnel need to know to properly care for you in an emergency.

☐ By checking this box and signing below, the employee indicates their preference to not share personal medical information with the National Park Service, Great Lakes Network Office. The employee understands that his/her refusal may delay or hinder necessary care if he/she has an undisclosed underlying medical condition or drug allergy. The employee is free to change their mind at any time and submit a completed form.

Employee

Date

Completion of this form is voluntary and does not influence any hiring decision or performance evaluation. It is strictly to be used in case of an emergency when the person named above is unable to provide such information to emergency personnel. This form will be kept in a locked file at the Network office and will be destroyed by the employee at the end of their term of employment.

Standard Operating Procedure #3: Using the Global Positioning System (GPS)

Version 1.1

Prepared by

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Revision History Log

The following table lists all edits and amendments to this document since the original publication date. Information entered in the log must be complete and concise. Users of this standard operating procedure will promptly notify the project manager and/or the Great Lakes Network (GLKN) data manager about recommended and required changes. The project manager must review and incorporate all changes, complete the revision history log, and change the date and version number on the title page and in the header of the document file. For complete instructions, please refer to Revising the Protocol, SOP #13.

Previous Version Number*	Revision Date	Author (with title and affiliation)	Location in Document and Concise Description of Revision	Reason for Change	New Version Number
1.0	1/28/2016	David VanderMeulen, Rick Damstra, GLKN	Updated the NPS contact author and authorship, removed references to specific versions of software, fixed a handful of grammatical errors throughout.	To clarify and update SOP based on changes that have occurred since Version 1.0 was published in 2008	1.1

*Add rows as needed for each change or set of changes tied to an updated version number

Acknowledgements

Major portions of this SOP have been adopted from the GLKN GPS Field Data Collection Guide. The guide was originally published as Appendix K to the Great Lakes Inventory and Monitoring Data Management Plan (Hart and Gafvert 2005). Appendix A of this SOP was also adopted from that document and contains a summary discussion of the general types of GPS units available, as well as definitions of terms commonly associated with use of GPS units.

3.0 Introduction

This Standard Operating Procedure (SOP) provides guidance on some of the more common operations associated with global positioning system (GPS) units. Although most GPS units are capable of multiple functions, only those operations relevant to the protocol established to monitor water quality on large rivers are discussed.

Water quality monitoring staff utilize both mapping and recreational-grade GPS units. The Trimble GeoXT is an example of a mapping-grade GPS unit, while the Garmin 76S is an example of a recreational-grade unit. Mapping-grade units record data with location accuracy ranging from sub-meter to less than five meters. Recreational-grade units generally are not as accurate, with spatial accuracy less than 15 meters. Mapping-grade units have greater ability to capture spatially referenced metadata (i.e., attributes) than recreational-grade GPS. A detailed discussion of these types of GPS can be found in Appendix A.

Software pertinent to the use of these GPS units includes:

- Trimble TerraSync mapping-grade GPS unit software
- Trimble GPS Pathfinder Office desktop GPS processing software
- ESRI ArcGIS desktop GIS software
- Minnesota Department of Natural Resources DNRGPS Application – desktop application to transfer data between recreational-grade GPS units and GIS software
- NPSTORET software, NPS database for water quality data (refer to SOP #8, Data Entry and Management).

Due to the rapid development of commercial software and hardware capabilities, it is likely that other GPS units or software will be utilized in the future. Therefore, this SOP is meant to act as a working document that is updated periodically as new hard-ware becomes available. Although nomenclature may differ depending on what hard-ware is utilized, this document should provide sufficient guidance on the general process of data collection using GPS tools until the SOP is revised. It is strongly recommended that water quality monitoring staff obtain unit-specific GPS training prior to deploying in the field. The training should include hands-on use, and should be designed to test all appropriate functions and operations prior to going out into the field.

Additionally, this SOP is not intended to be exhaustive or simply a regurgitation of operating manuals, but a document that might be carried into the field or periodically reviewed by field technicians and project leaders. Although some of the information in this SOP is specific to the water quality monitoring effort by GLKN, much of the text contained herein is applicable to other data collection efforts by GLKN staff. The objective of this document is to summarize GPS use guidelines applicable to the water quality monitoring efforts for large rivers at GLKN parks.

3.1 Role of GPS in Water Quality Monitoring on Large Rivers

GPS units are primarily used in the following ways to support water quality monitoring on large rivers:

1. For water quality monitoring on large rivers, navigating to and sampling at the position locations of monitoring stations is accomplished through a combination of:
 - Using GPS units to navigate to previously established GPS point locations
 - Using hardcopy topographic maps and written notes with site descriptions
 - Landmarks, such as specific posts on a bridge site, or other distinctive landscape features
 - Comparing digital photos taken during site establishment with observer ocular estimates
 - Field experience of NPS staff conducting the monitoring
2. Accurate station location descriptions must be recorded and carefully followed by sampling personnel on subsequent field visits for water quality monitoring. Therefore, once on station, GPS equipment is used to obtain the station's coordinates, which allows the user to spatially reference water quality monitoring data to specific geo-referenced locations for each sampling event.

Additional uses of the GPS units may develop over time if protocols change or other needs are identified. For example, the Great Lakes Network currently promotes the use of electronic data logging equipment to enhance data quality and simplify data management. Many water quality parameters (e.g., pH) are now primarily recorded with electronic data logging equipment, while others (Secchi disk depth, names of observers, etc.) are logged on hardcopy only. Therefore, GPS units capable of logging more information other than just waypoints (e.g., Trimble GeoXT) could be programmed to allow for electronic recording of these parameters. The next two sections of this SOP will outline some of the more common and pertinent functions associated with mapping and recreational-grade GPS units, which are the types of units currently used by GLKN staff for water quality monitoring.

3.2 Using Mapping-Grade GPS Units

All mapping-grade GPS users should become familiar with GLKN GPS collection procedures and relevant manufacturer's user guides and operating manuals before GPS operation. For example, prior to using a Trimble GeoXT (mapping-grade) GPS unit, the following documents should be reviewed:

- Appendix A of this SOP
- GeoExplorer CE Series: Getting Started Guide
- GPS Mapping for GIS with TerraSync and GeoExplorer CE Series or TerraSync Operation Guide v2.4x

- Basic GPS Data Capture Using TerraSync: A Quick Start Guide

Mapping-grade GPS units provide the user with a variety of tools for field data collection. GLKN encourages the use of these units for most projects. These units can be used to acquire spatial data related to points, lines, and polygons along with associated, user defined, tabular attributes. Careful forethought and advanced planning are required to take advantage of these capabilities long before data collection begins.

3.2.1 Data Dictionaries

TerraSync software on mapping-grade GPS units is capable of using data dictionaries. Data dictionaries define the structure and rules to store attribute information about the feature being mapped and are customized for each project. GLKN data management personnel should be directly involved in the creation of data dictionaries. Basic steps include:

1. Identify the features to be mapped. These features are real world physical locations of objects (e.g., a water quality monitoring station) that are categorized as point, line, or polygon features.
2. Identify the attributes to be collected for each feature while in the field and create a data dictionary. Part of this process is assigning a unique identifier to each feature. (For example, if a survey plot is mapped as both a point and a polygon, one feature should be named plotname_poly and the other plotname_pt.).
3. Implement and test the data dictionary. Field staff should conduct a complete trial run for newly-created data dictionaries before beginning field work. Corrections and refinements are inevitable after such a trial.

Because most of the data for water quality monitoring on large rivers are recorded by electronic sampling equipment, there has not been much incentive to utilize data dictionaries. However, as discussed earlier, it is possible that the few parameters not currently logged electronically could be incorporated into a data dictionary and recorded. As stated above, it is highly recommended that GLKN data management personnel be involved in this process, and that once created, the data dictionary only be altered with permission from data management personnel. This will reduce the potential for confusion and mistakes when data are processed at the end of the field season.

3.2.2 GPS Settings

Positional accuracy of GPS data can be affected by several factors that can be monitored and recorded with mapping-grade GPS units. Table 3-1 lists these factors, their definitions, and the standard settings for GLKN field work. All spatial data collected shall be analyzed for spatial accuracy and shall meet or exceed the National Map Accuracy Standards (Table 3-1 in Appendix A, and <http://mapping.usgs.gov/standards/>). Table 3-2 indicates the coordinate system settings for data collection in GLKN parks.

Table 3-1. GPS receiver settings, definitions, and standards for use at GLKN parks.

Setting Name	Definition	GLKN Setting Standard
Almanac	File containing estimated position of satellites, time corrections, and atmospheric delay parameters	Acquired automatically by GPS unit or from online sources within 10 days prior to GPS field work
Altitude reference	Ellipsoid model	Height above Ellipsoid (HAE) (preferred) or Mean Sea Level: if MSL is used, indicate Geoid Model
Antenna height	GPS antenna height above the ground	Variable, usually 1.0 meters for handheld and 1.5 m for backpack
Datum	Geodetic model designed to fit a point on the earth's surface to an ellipsoid	NAD 83 (CONUS) [preferred] WGS 84 [GPS default, as fallback] NAD 83 (CONUS) (CORS 96) [for H-Star use with GeoXH Trimble GPS Unit]
Elevation mask	The minimum angle above the horizon at which a GPS receiver will track a satellite	15 degrees
Feature types	Geometry of spatial data	GIS native formats; point, line and polygon are preferred
Logging interval	Time interval between the recording of individual GPS fixes	Points: 1 second Lines and Polygons: 5 seconds, but 1 second in some circumstances
Minimum fixes for point positions	Number of GPS fixes that are used to calculate a single position for a point feature	50 fixes (120 points in the case of GeoXH)
Mode	2 dimensional for horizontal positions and 3 dimension with an elevation position	3-dimensional (4 satellite minimum)
PDOP mask	Positional Dilution of Precision, a GPS quality estimate based on satellite geometry	6.0 or less
Real-time settings	GPS unit may be capable of performing differential correction of data during collection	Select Integrated WAAS (unless using H-star, e.g., XH unit); setting will be 'auto' or 'on'
Satellite vehicles	Number of satellites used for position fixes	4 minimum
SNR mask	Signal-to-Noise ratio is a measure of the satellite signal relative to background noise	4.0 minimum, 6.0 or greater preferred
Unit of measure	Linear unit of measure	Meter (metric)

GPS signals are received in the WGS84 datum. Processing and transformation of the positional information to other datums can take place internally in the GPS unit or in software, either the GPS data processing software (see below) or in GIS software. GPS data that will receive no post-

processing differential correction, unlikely with a mapping grade receiver, can be collected in native WGS84 or NAD83 (CONUS), and ensuring that the datum used is recorded in the metadata. More likely with a mapping grade receiver, the GPS data will be differentially corrected after collection using data from one or more reference base stations. Data on the GPS unit should be collected in the same datum as the data are output from the reference base stations. The majority of public base stations in the US are part of the National Geodetic Survey's Continuously Operating Reference Station (CORS) network; and output information in the NAD83 (CONUS) CORS datum. Setting the GPS unit to record data in NAD83 (CONUS) CORS datum will result in the most accurate spatial information when differentially corrected against a CORS base station. A very high precision mapping grade unit may use H-Star technology, which requires post collection differential correction; using the NAD83 (CONUS) CORS datum and no real-time correction (WAAS) on the GPS unit, and correcting against a group of CORS base stations is necessary to realize the maximum accuracy from an H-Star receiver.

Table 3-2. Coordinate system settings for Great Lakes Network parks. **Bold font** indicates parks where water quality monitoring on large rivers will take place.

Park	UTM Zone	Datum
APIS	15	NAD 1983 (Conus)
GRPO	16	NAD 1983 (Conus)
INDU	16	NAD 1983 (Conus)
ISRO	16	NAD 1983 (Conus)
PIRO	16	NAD 1983 (Conus)
MISS*	15	NAD 1983 (Conus)
SACN*	15	NAD 1983 (Conus)
SLBE	16	NAD 1983 (Conus)
VOYA	15	NAD 1983 (Conus)

* Indicates parks where water quality monitoring on large rivers will take place (also in bold font).

Before beginning data collection, the GLKN data management staff or the GPS user should complete some mission planning tasks. If high accuracy fixes are desired and there is some flexibility in scheduling of a field data collection mission, the user should complete a satellite survey to determine the best timing of the mission, usually when the most satellites are visible and in the best geometry. Software, such as Trimble's Quick Plan (also included in Pathfinder Office) can be used to look for time windows that should offer the lowest PDOP readings, and thus the highest positional accuracy.

If a project requires navigation to preset locations or waypoints, this information must be pre-loaded onto the GPS hardware before starting a field data collection mission, unless the locations have been previously stored on the GPS. Having printed copies of bathymetric maps with sampling sites marked is a good backup and can maximize field time and efficiency. Some of the sampling stations on large rivers are located relative to unique features (e.g., bridges), which are annotated on

topographic maps. Detailed descriptions and photographs of the shoreline are also used to navigate to designated sampling sites.

Many mapping-grade GPS units have the capability of storing and displaying background maps or GIS layers, which can be helpful when navigating in the field. For example, topographic maps of the St. Croix River can be used with GPS units to find water quality sampling stations. The GLKN data management staff can support preparation of these background layers and, if necessary, assist in loading them onto the GPS hardware.

GPS units create files to store data during a field session using a prefix and date-time stamp as file names. For example:

RMMDDHHx

R – Unit Prefix

MM – Month

DD – Day

HH – Hour

X – a, b, c, etc., the order files are created within an hour

If multiple GPS units are used for a project, a unique prefix (letter) should be assigned to each unit, which will ensure that downloaded files for each unit contain a unique identifier within the filename. For example, with three GPS units, the unique letters for the units could be N, G, and A. Those letters would serve as a prefix for the file n (e.g., N102715A, G102715A, and A102715A would indicate units N, G, and A, October 27, 15 hour, A first in hour). In addition, if a data dictionary is used, and entry of the observer's name or initials is not an option in the dictionary, then the observer's name or initials should be included in the file name.

Each user should be familiar with the capabilities of the GPS hardware and field computers. If possible, water quality monitoring staff should receive hands-on instruction from someone familiar with the equipment. At a minimum, the equipment user guides and operator's manuals should be reviewed, and the operator should test the functions s/he intends to use.

It is extremely important that each user become familiar with the battery power and memory capabilities of the GPS units. All units have limited battery and memory resources; these features should be thoroughly tested to gain an understanding of the power and memory limitations of the GPS units before being deployed in the field. It is possible to power or recharge these units from a DC power source, such as a vehicle power outlet.

3.2.3 Data Collection

Data collection should be performed using an approved data dictionary or database for the protocol, if applicable. Moreover, users should be mindful of the following concepts:

- If using a GPS unit that gives an approximate 5m horizontal accuracy, the user cannot map anything as a polygon that is less than 4 to 6m in width or diameter. Such objects must be captured as point features.
- If a GPS user is collecting a line or polygon feature and then stops moving, the GPS unit will continue to collect data (Figure 3-1, examples 1 and 2). Users need to be familiar with the Pause/Resume toggle key and use it liberally. This technique greatly improves subsequent data quality and reduces the need for time-consuming spatial editing.
- Another way to avoid errors is to collect point features that represent the beginning and end points of a line transect (Figure 3-1, examples 3 and 4). Having these reference point locations will mean easier editing of any zig-zagging line features.

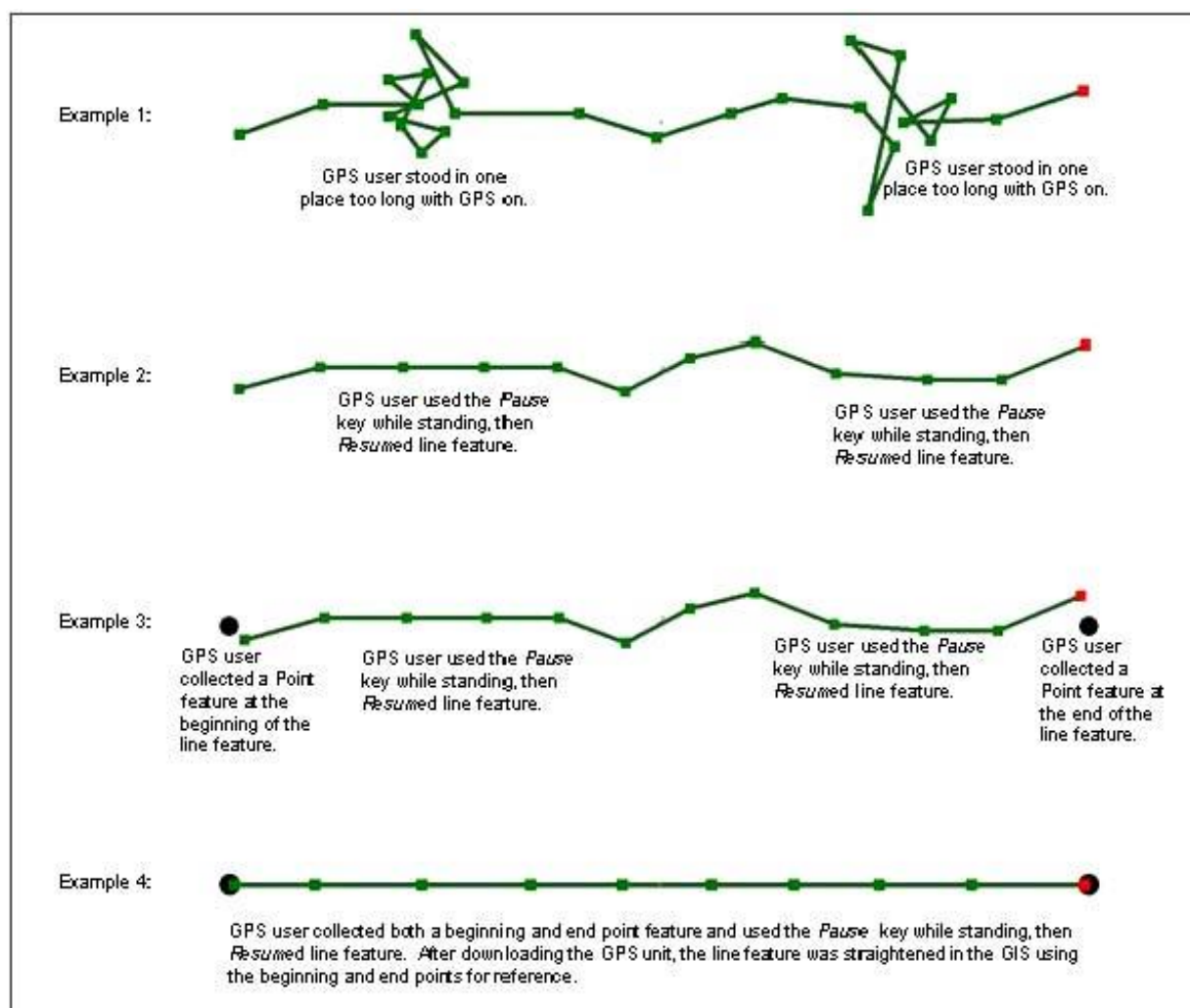


Figure 3-1. Examples of GPS point and line features data collection.

Mapping-grade GPS units have additional features that aid in data collection. These include:

- *Nested Features*—Allows user to collect a point while collecting a line or polygon feature. For example, while surveying potential amphibian habitat along a stream (line feature), the user can pause the line feature, take a point for a specific observation, then resume the line feature collection.
- *Offset Feature*—Allows user to collect a feature when topography is such that getting next to or over the feature is impossible. For example, a GPS line could be collected while taking a horizontal zooplankton tow off the side of a boat.
- *Between-feature Positions*—the GPS unit collects positions without any feature or attribute data. This feature is useful for tracking areas traveled during a day. For example, while surveying for presence or absence of invasive plants, a user could collect data on the area surveyed in addition to locations of specific plants found.

3.2.4 Data Processing

When data collection is complete for the day or round of sampling, data are downloaded from the GPS unit to a computer. For Trimble GPS units, the proprietary software Pathfinder Office (or the GPS Analyst extension) is used to download, differentially correct, and then export the data to a GIS format. [Note: Trimble Pathfinder Office and GPS Analyst are relatively expensive. However, Trimble also offers a free data transfer utility to download data from the GPS units to a Windows-based PC]. Differential correction is a post-processing procedure to improve upon raw GPS positions using base station data. Base stations consist of a GPS antenna and receiver positioned at a known location specifically to collect data from satellites. The distance between the base station(s) and the remote GPS receiver should be kept to a minimum.

Differential correction should be conducted on all GPS data collected, even if data were collected using the real-time collection feature. Once the data are differentially corrected, they can be verified and edited. Unintentional features can be deleted and attributes can be reviewed.

The last step in processing data is exporting the data set to GIS (ArcGIS or ArcView). Depending on the software used for this process, newly created files generated when exporting data are often assigned generic names. For example, if Pathfinder Office is used to export a file named 'SACN2007.cor' (.cor denotes that the file has already been differentially corrected) that only contains point features, the exported file will be named 'point_ge.shp'. Great care should be taken to not overwrite this file when exporting other data, as the software will continue to use this generic naming convention the next time it is used. In addition, during the export process, the coordinate system to which the data will be exported to should be verified (Table 3-2).

Additional data attributes can be included in the data exports. Data attributes recommended by GLKN are listed in Table 3-3.

Table 3-3. Recommended fields to be exported in addition to GPS features.

All Features	Point Features	Line Features	Area Features
PDOP	Height	Length (2D)	Area (2D)
Correction status	Position	Length (3D)	Perimeter (2D)
Receiver type	--	--	Perimeter (3D)
Date recorded	--	--	--
Data file name	--	--	--
Total positions	--	--	--
Data dictionary name	--	--	--

Managing the incoming GPS data can be a challenge, especially if there are multiple units per project. Common practices used by GLKN include:

- Download all data to a computer or network drive that is regularly backed up.
- Keep GPS data and GIS data separate through electronic file management.
- Directories and files names should not contain non-alpha-numeric characters and/or spaces (except underscores).
- Keep GPS data in well-organized directories (see Hart and Gafvert (2005), GLKN Data Management Plan, for more details).

At the end of a project, all data and background files should be removed from the GPS unit to free available memory. Data files should not be left on a unit if they have been properly downloaded and verified. In addition, some GPS units require their batteries to be re-charged periodically. Failure to do so can cause the GPS unit batteries to discharge completely, and may cause some files and software to be deleted.

Additional information can be found at www.nps.gov/gis/gps/gps4gis/, which describes the steps outlined here in greater detail.

3.3 Using Recreational-Grade GPS Units

Recreational-grade GPS units can be used to acquire location information (generally points) when spatial accuracy is not paramount to the project. Recreational GPS units do not have data dictionaries for storing attribute information with the point location. However, using a recreational-grade unit to capture a waypoint at each sampling site is a reliable means to verify the correct sampling site has been reached, even if a GPS location is not needed.

As with mapping-grade GPS units, personnel that employ recreational-grade GPS units should become familiar with GLKN GPS collection procedures and relevant manufacturer's user guides and

operating manuals before GPS operation. For example, prior to using a Garmin 76S (recreational-grade) GPS unit, the following documents should be reviewed:

- Appendix A of this SOP
- GPSMAP 76S Quick Start Guide
- GPSMAP 76S Owner's Manual and Reference Guide
- Garmin MapSource™ User's Manual and Reference Guide

3.3.1 Planning

If a recreational-grade GPS meets the criteria of the project, the unit chosen must have the capability of downloading collected data to a personal computer. Downloading data is usually accomplished with a parallel or USB cable connection.

Much of the data collected by GPS will eventually reside in a relational database. Each GPS feature collected should contain a unique identifier that relates the feature to an associated record in a database. For water quality monitoring on large rivers, the records associated with each GPS feature will consist primarily of water quality parameters. Since recreational GPS units have only one text field for input, careful consideration should be given to the use of this field and the design of unique identifiers. GLKN data management and GIS staff can assist in creating unique IDs on a project-by-project basis.

3.3.2 Data Collection

Location data are captured by recreational-grade GPS units as *waypoints*. When taking a waypoint, enter the site ID or site designation in the text field provided. It is also good practice to collect reference points at regular intervals. These reference point positions should be taken at known locations (e.g., trailheads, parking lots, stream confluences) which can later be used in GIS to check the accuracy of waypoint data.

If navigation to preset waypoints is applicable to a project, the waypoints must be loaded onto the GPS unit before departure to the field. Some recreational grade GPS units have the ability to store and display topographic maps, which can aid in navigation. Printed topographic maps of the waypoint locations can also be used to maximize field time and efficiently navigate between waypoints.

3.3.3 Data Processing

Data should be downloaded from GPS units once a day or after each field session. The DNR Garmin freeware product (www.dnr.state.mn.us/mis/gis/tools/arcview/index.html) can be used to download data from Garmin GPS units. Data should be downloaded both as a text file and a shapefile. Each file name should include the download date. Points should be checked for reasonable spatial accuracy and errors. Subsequent downloads should be error-checked in the same manner. When data collection is finished, all files should be compiled into one spatial file, and along with the raw downloads, should be saved to the appropriate location on Great Lakes Network servers (refer to SOP #8, Data Entry and Management for more detail).

3.4 Metadata

Regardless of the type of GPS unit used to collect data, all resulting GIS datasets need to have information documenting how the GPS data were collected. NPS requires that FGDC (Federal Geographic Data Committee, www.fgdc.gov/index.html) compliant metadata be written for all geospatial layers created (Executive Order 12906).

Until final FGDC metadata is written, the data collection and management process is incomplete. Tracking GPS projects depends on the complexity of the project, how many participants, length of project etc. Documentation can be a simple 'readme' text file, or a detailed daily log.

The Great Lakes Network recommends formal metadata be written by the data collectors, as they are the ones familiar with the project and resulting data. However, Network data management and GIS staff usually end up documenting someone else's work. Chapter 7 of the GLKN Data Management Plan (Hart and Gafvert 2005) includes a detailed discussion of metadata procedures. At a minimum, the following details should be documented to facilitate final FGDC metadata:

- Name of project
- Name(s) of data collectors
- EHE/EPE or maximum PDOP (using 4 satellites)
- Coordinate system (projection, datum, and zone)
- Type (or types) of GPS units used
- The range of field collection dates
- Name of base station(s) used for differential correction
- Name and version of software used for downloading
- Any major editing performed on the raw data (e.g., moving of points)
- All versions of data dictionaries used

3.5 GPS and NPSTORET

Water quality data, including chemical, physical, and biological data, are managed according to guidelines from the NPS Water Resources Division. These guidelines include using the NPSTORET desktop database application to help manage data entry, documentation, and transfer. The Network oversees the use of NPSTORET according to the Network's water quality monitoring protocols and ensures the content is transferred at least annually to NPS Water Resource Division for upload to the EPA STORET (STORage and RETrieval) database.

NPSTORET requires that every water quality monitoring station location must have an assigned latitude and longitude coordinate. Also, the horizontal datum to which these coordinates are referenced (typically North American Datum 1983 or World Geodetic System 1984) and the method by which they were obtained (GPS, map interpolation, etc.) must be provided. Therefore, the GLKN

project leader for monitoring water quality on large rivers and the GLKN data manager collaborate in organizing monitoring station coordinates in an acceptable format (e.g., Excel spreadsheet or Access database) to be imported to the NPSTORET database. A detailed discussion on using the NPSTORET database is found in SOP #8, Data Entry and Management.

3.6 QA/QC

Long-term monitoring is only useful if stakeholders have confidence in the data. Efforts to detect trends and patterns in ecosystem processes require high-quality, well-documented data that minimize error and bias. Data of inconsistent or poor quality can result in loss of sensitivity and lead to incorrect interpretations and conclusions.

NPS Director's Order #11B: Ensuring Quality of Information Disseminated by the National Park Service (www.nps.gov/policy/DOrders/11B-final.htm) specifies that information produced by the NPS must be of the highest quality and based on reliable data sources that are accurate, timely, and representative of the most current information available. Therefore, GLKN will establish and document procedures for quality assurance (QA) and quality control (QC) to identify and reduce the frequency and significance of errors at all stages in the data life cycle (see SOPs #8 and #12 for details on data management and QA/QC, respectively). Under these procedures, the progression from raw data to verified data to validated data implies increasing confidence in the quality of those data. Quality assurance and quality control procedures will document internal and external review processes and include guidance for addressing problems with data quality.

Examples of general of QA/QC practices include:

- Standardized field data collection forms
- Use of field computers and automated data loggers
- Proper calibration and maintenance of equipment
- Training of field crew and data technicians
- Database features such as built-in pick lists and range limits to reduce data entry errors
- Automated error-checking routines

Many of the standard operating procedures associated with the protocol for monitoring water quality in large rivers include a discussion of QA/QC as it relates to the protocol. Examples of QA/QC practices pertaining to use of GPS include:

- Ensure that GPS-related software is periodically updated as it becomes available and has been tested.
- Record location positions on field data forms as well as with the GPS unit.
- For each monitoring station, compare location positions for different sampling events, including the position recorded during establishment of the monitoring station. This will allow for an assessment of position accuracy over time.

- If data dictionaries are used with mapping-grade GPS units, ensure that the coordinates for the monitoring station match the other attributes recorded on the GPS unit for that monitoring station.
- Check to see if the accuracy of the GPS unit meets or exceeds the National Map Accuracy Standards shown in Table 1 of Appendix A.
- Ensure that the appropriate coordinate system is used when collecting and exporting data.
- Use mapping software (e.g., Pathfinder Office or ArcGIS 9.x) to view waypoints (or features) overlaid on a geo-referenced air photo or topographical map to check for accuracy.
- If applicable, check the accuracy of the attribute(s) recorded on a GPS unit by using mapping software (see bullet above) and look-up tables or in spreadsheets generated after post-processing is complete.

A final report on data quality, including data collected by GPS, will be incorporated into the documentation for this project. Such documentation will include a listing of the specific methods used to assess data quality and an assessment of overall data quality prepared by the project manager. This is a necessary part of the data quality elements of the metadata file.

3.7 Literature Cited

Hart, M., and U. Gafvert (editors). 2005. Data management plan: Great Lakes Inventory and Monitoring Network. National Park Service Great Lakes Inventory and Monitoring Network Report GLKN/2005/20. National Park Service, Ashland, Wisconsin.

Appendix 3A

This appendix discusses the role of GPS in GLKN data management and provides an explanation of the types of GPS units that are available. The text is taken directly from the GLKN GPS Field Collection Guide, which can be found in its entirety in Appendix K of the Great Lakes Inventory and Monitoring Data Management Plan (Hart and Gafvert 2005).

Introduction

Over the past decade new tools have been developed to help researchers collect and manipulate data while in the field. Global Positioning System (GPS) is one such tool. GPS is currently a constellation of 28 US Department of Defense satellites (as of 2006) orbiting 11,000 miles above the Earth, making a complete orbit approximately every 12 hours, and transmitting signals to Earth at precisely the same time. The position and time information transmitted by these satellites is used by a GPS receiver to triangulate a location coordinate on the earth using three or more satellites.

Role of GPS in GLKN Data Management

Data collected using GPS-enabled equipment represents all or part of the acquisition stage of an information resources lifecycle that includes several other stages (see Section 5.4 in GLKN Data Management Plan). The process and methodology used for acquisition, planning, data collecting, and post-processing incorporate several aspects of data management, including quality assurance, data storage and organization, and data stewardship. To promote data quality and simplify data management, the Great Lakes Network expects to use electronic data logging equipment for some data acquisition. However, parallel or complementary use of hand written data sheets and field notes will remain important for data collection activities.

Types of GPS Units

At the most basic level GPS equipment can just consist of a GPS antenna and the associated signal processing circuitry. The antenna can be a standalone device, be incorporated in to a handheld unit, or be integrated into a larger electronic device, such as a personal data assistant (PDA), data logger, or portable computer. As technology evolves, the Great Lakes Network will continually try to use equipment which maximizes spatial accuracy; reduces hardware weight and user fatigue; and reduces database development, data manipulation, and transformation.

There are three major types of GPS units that are based on the level of accuracy to which spatial data can be collected. Survey-grade GPS units are used for surveying tasks that require very high accuracy (1 cm or less). Mapping-grade units can map features from sub-meter to less than 5m accuracy, employing differential correction. Recreational-grade GPS units are sold primarily for outdoor sports and recreational activities. Accuracy using recreational GPS units ranges from 5 to 30m. Most natural resource-related data collection requirements correspond to either the recreational-grade or mapping-grade. Figure 3-A1 shows some of the major differences between these two types.

Deciding which type of unit to use is an essential part of project planning, and depends on the end product needed. Mapping-grade GPS units are recommended for most GLKN field work; however, for some projects recreational-grade units can meet a project's accuracy requirements and reduce the

cost of field operations. The choice of GPS unit should be made by the project manager after consulting with the GLKN data management and GIS staff.

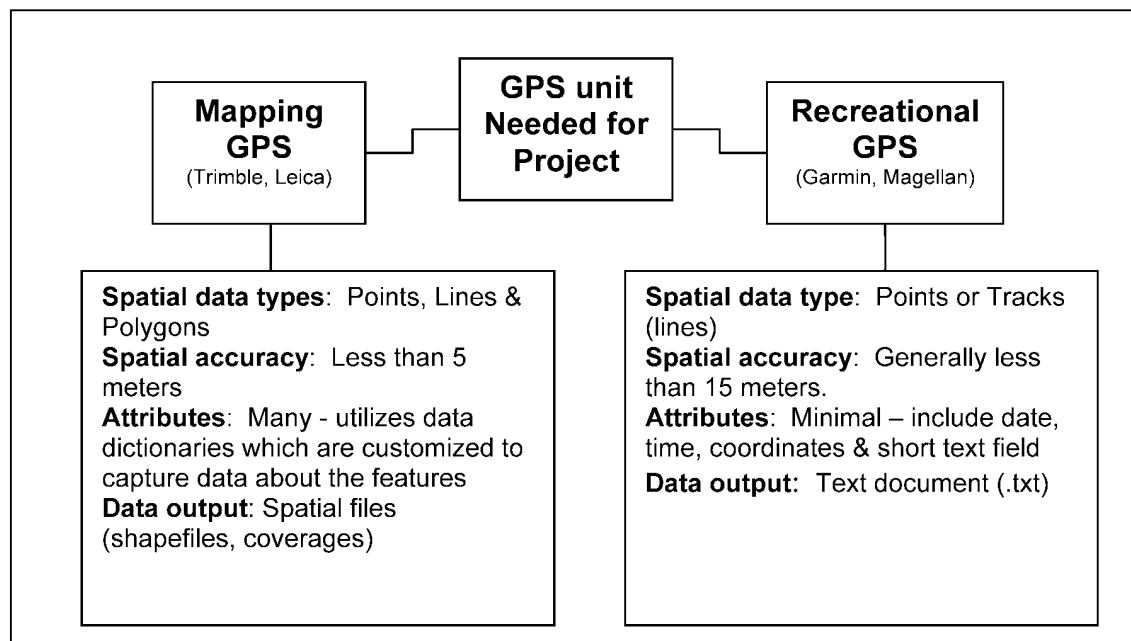


Figure 3-A1. Differences between different grades of GPS units.

All resulting GIS data layers need to meet or exceed the National Map Accuracy Standards for a 1:24,000 product (NPS GIS Data Standards, 2002, www.nps.gov/gis/data_info/standards.html). Table 3-A1 provides the allowable horizontal accuracy for some common scales.

Table 3-A1. Map scales and allowable error.

Scale	Allowable Error
1:40,000	33.8 meters (111 feet)
1:31,680	16.1 meters (53 feet)
1:24,000	12.2 meters (40 feet)
1:20,000	10.1 meters (33 feet)
1:12,000	6.1 meters (20 feet)
1:9,600	4.9 meters (16 feet)
1:4,800	2.4 meters (8 feet)
1:2,400	1.2 meters (4 feet)
1:1,200	0.6 meters (2 feet)

Definitions

Accuracy: The degree of conformance between the estimated or measured position, time, and/or velocity of a GPS receiver and its true time, position, and/or velocity as compared with a constant standard.

Almanac: Data transmitted by a GPS satellite, which include orbit information on all the satellites, clock correction, and atmospheric delay parameters. The almanac is used to facilitate rapid satellite vehicle (SV) acquisition. The orbit information is a subset of the ephemeris data with reduced precision.

Attribute: Tabular information about a specific feature.

Base Station: GPS files collected continuously from community base stations, local base stations, or Continually Operating Reference Stations (CORS). Gathering base files will require an internet connection and software that dials into a server that houses the base station data collected at the same time of the rover. Data stored on these servers will not be available in real-time - hence this step is conducted after field collection. Trimble users would use the Differential Correction utility supplied in Pathfinder Office.

Differential Correction: The merging of rover file data with base map data to correct position errors due to atmospheric interference. Autonomous data (rover) are collected in the field while base data are stored at the stationary base station. The two datasets are loaded into a post-processing software package where corrections are applied. This process will reduce errors in the field collected data (the rover) by correlating and correcting for known errors recorded in the base file that has the same time tag. As distance between the rover and base file increase, there is degradation in post-processed accuracy. In general, a degradation of one part per million (1ppm) occurs as the distance between the base station and rover increases. For example, one millimeter of degradation occurs for every kilometer between base and rover.

Datum (*geodetic datum*): A mathematical model that is designed to fit a point on the earth's surface to an ellipsoid. Commonly used datums are North American Datum (NAD) 1927, and NAD 1983, modeled to represent the North American continent.

Feature: A feature is the spatial location of a physical object, or some event or phenomenon. Features are often referred to as graphic data in a GIS. Examples include a tree (point), road (line), or land parcel (polygon).

FGDC: The Federal Geographic Data Committee is a 19 member interagency committee composed of representatives from the Executive Office of the President, Cabinet-level and independent agencies who develop policies, standards, and procedures for organizations to cooperatively produce and share geographic data. (www.fgdc.gov/index.html).

Global Positioning Systems (GPS): A constellation of a minimum of twenty-four satellite vehicles orbiting the earth approximately every twelve hours at an approximate pacing of sixty degrees, between 11,000 – 12,000 miles above the surface of the Earth

Lines: Geographic term related to the scale that describe how a feature is drawn. Lines are linear measures of a feature (such as a line representing a trail)

Mapping Grade: GPS receivers capable of attaining five meters of accuracy or better using differential correction.

Metadata: Data about the data. Usually comes in the form of a text or html document with information on the dataset's quality, current projection, attributes, distribution and citation. In the National Park Service, this generally implies a file compliant to the FGDC Content Standard for Digital Geospatial Metadata.

Multipath: Error which occurs when a GPS signal sent from a satellite vehicle is bounced or redirected by an object, prior to reaching a GPS receiver. Multipath will cause the time it takes a GPS signal sent by a satellite vehicle to reach a GPS receiver to be inflated. This will cause inaccuracies in positions collected.

Points: Geographic term related to the scale that describe how a feature is drawn. Points are single dimensional features (such as a point representing a spring).

Polygons: Geographic term related to the scale that describe how a feature is drawn. Polygons have area associated with the feature (such as a circle representing a parking lot).

Projection: A method of representing the earth's three-dimensional surface as a flat two-dimensional surface. This normally involves a mathematical model that transforms the locations of features on the earth's surface to locations on a two-dimensional surface.

Post Processing: Utilizing base station data, GPS software, and data acquired by a GPS receiver in the field to gain an accurate fixed position.

Triangulation: The process of determining the distance between points on the earth's surface by dividing up a large area into a series of connected triangles, measuring a base line between two points, and then locating a third point by computing both the size of the angles made by lines from this point to each end of the base line and the lengths of these lines.

Waypoint: A named three-dimensional position on the earth's surface, that is, having both a latitude and longitude. Waypoints are assigned to a fixed location in the field so it can be navigated to consistently and accurately through time.

Standard Operating Procedure #4: Site Establishment

Version 1.1

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Revision History Log

The following table lists all edits and amendments to this document since the original publication date. Information entered in the log must be complete and concise. Users of this standard operating procedure will promptly notify the project manager and/or the Great Lakes Network (GLKN) data manager about recommended and required changes. The project manager must review and incorporate all changes, complete the revision history log, and change the date and version number on the title page and in the header of the document file. For complete instructions, please refer to Revising the Protocol, SOP #13.

Previous Version #	Revision Date	Author (with title and affiliation)	Location in Document and Concise Description of Revision	Reason for Change	New Version #
1.0	2/10/2016	Rick Damstra and David Vandermeulen, GLKN	Updated, NPS contact author and authorship, removed reference to calculation of stream discharge and rating curves. Also removed reference to installing permanent site markers and staff gauges.	Not practical or necessary for site establishment, in current park units.	1.1

*Add rows as needed for each change or set of changes tied to an updated version number

Acknowledgements

The authors would like to acknowledge Susan O’Ney, who blazed the trail when it came to writing standard operating procedures for National Park Service water quality monitoring protocols. Much of our site establishment SOP is based on her work. Thanks also go to Pete Penoyer for his guidance.

4.0 Introduction

Research on potential sites and preliminary site visits are required to formally establish the specific monitoring stations. Information on the river cross-section, depth, and flow velocity must be gathered, and discharge rating curves for sites where gauging stations are nearby should be obtained. Detailed information on each sampling site must be compiled into field binders and must include all material necessary for safe and efficient field operations.

4.1 Establishing NPSTORET Project Files

Project files should be established prior to site reconnaissance visits using the NPSTORET database template. The project manager is responsible for ensuring electronic and paper project and site files are established and maintained. NPS policy requires specific information on surface water sampling sites. Refer to Tucker (2004) for detailed discussions of NPS data and metadata requirements, and to SOP #8 of this protocol for detailed instructions on data entry and management.

4.2 Selection of Proposed Monitoring Stations

The site selection process and proposed monitoring stations for monitoring water quality in large rivers are described in Section 2 of the protocol narrative. The selection process includes oversampling of the randomly-selected sites; if any of the proposed randomly-selected monitoring sites prove to be inappropriate or not representative of the resource being monitored, than a replacement must be selected from the pool of oversampled sites.

4.3 Permission for Research

Once a proposed station has been identified, permission should be obtained from the appropriate park to conduct research and establish permanent markers within park boundaries. Contact the park's research coordinator and complete all required permit application forms prior to conducting site establishment activities.

4.4 Location Reconnaissance and Assessment

Once a potential monitoring station has been selected and its GPS location verified on a map, the site should be visited to assess whether it is an appropriate monitoring location. Water quality monitoring sites and flow monitoring sites for a given monitoring station do not need to be at the same exact location, but they should be within one mile of each other and represent the same flow conditions (no sinks or sources between the two sites). Key concerns are accessibility, representativeness, and stability.

4.4.1 Accessibility

Bridge crossings provide safe access to large rivers during high flows, but can be the source of road contaminants (volatile organic compounds and especially road salts), and rarely cross at randomly-selected locations. Therefore it is preferable to sample from boats. Identify the nearest public access boat landing. The landing should be located within a few miles (<10 miles) of proposed monitoring

site. Visit the site using the boat that is planned for field work to verify that average flow conditions are passable using that boat and the route is not blocked by riffles or rapids. Select monitoring locations that allow safe sampling during high flow with minimal risk to sampling personnel.

4.4.2 Representative of Average Flow and Water Quality of River Reach

At the proposed monitoring location, river discharge should be free-flowing (not stagnant) but not overly aerated or churned. The flow conditions should represent well-mixed but laminar flow. As part of the reconnaissance, verify that a proposed site is not immediately downstream of a pollution point source. If a site has been selected randomly to represent the average conditions of the river, then it should be located at least five stream-widths downstream from a tributary to ensure complete mixing of waters (Stednick and Gilbert 1998). During reconnaissance of a potential site, verify vertical and lateral mixing, as measured by core suite readings of the stream cross-section (see section 4.6 below).

4.4.3 Stable Setting for Long-Term Monitoring

The proposed monitoring site should be located along a straight portion of the river channel; avoid meander bends or side channels that may experience changes in the streambed morphology. It is best if the site is entrenched between stable river banks; avoid low banks or levees on one or both sides of the river that are prone to overbank flow, unless this is representative of the average conditions for this reach. For discharge measurements, it is preferable that the selected location is confined between bedrock or high clay banks; avoid sandy streambeds, if possible.

4.5 Permanent Marker and Site Coordinates

Accurate station location descriptions must be recorded and followed carefully by sampling personnel on subsequent field visits. Use Global Positioning System (GPS) equipment to obtain the station's coordinates and elevation near the water's edge. Read and follow SOP #3 for more details on using a GPS unit.

4.6 Photographic Documentation

Each sampling site must be documented with a series of photographs to establish site conditions. These photographs will be used to accurately relocate monitoring sites and to document field conditions. Document initial conditions at the site: take digital photographs of the site from a landmark on shore (e.g., boat landing, bridge, campsite), and photographs of the river from the site in the upstream and downstream directions.

Photographs should be identified with:

- The photograph number
- Date and time, even if the photographs are automatically date stamped by the camera
- The subject
- The location in narrative format and GPS coordinates
- The photographer

- Witnesses, if any
- The location from which the photograph was taken
- A short narrative description of the photographs

New photographs should be taken at the beginning of each field season or whenever site conditions change substantially.

4.7 Creating Field and Office Binders

Selected information that is needed for reference while at a sampling station is kept in a field binder. The field binder contains information used by trained personnel to locate and safely collect water quality samples and must be taken along on each sampling trip. Assemble and update the field binders with information such as description of sampling station, maps, data collected at station previously, and sampling instructions.

The office binder should contain reference information relevant to general field sampling procedures with tabs identifying each procedure for easy access during field work, including QA/QC reminders, a copy of the protocol narrative and all SOPs, copies of equipment instructions and troubleshooting; extra field forms, material safety data sheets for field supplies that contain hazardous chemicals or materials, and analytical service request and chain of custody forms.

See SOP #1, Pre-season Preparation, for more detailed information on preparing field and office binders.

4.8 Enter Station Information into NPSTORET

Upon returning to the office, enter all information into NPSTORET following the guidelines for establishing a new station record in GLKN's data management plan (Hart and Gafvert 2005). Station IDs should be assigned to new stations based on park unit, river name, and river miles (to the nearest 1.0 mile). Prepare a description of the location and compile other required information (including digital photographs) for establishing NPSTORET files.

In addition, information related to the water quality measurements collected at each site must be entered into NPSTORET. For a complete list of water quality variables and relevant metadata (i.e., analytical methods), see SOP #7, Samples and Analytical Laboratory Requirements. Follow details for data entry provided in SOP #8, Data Entry and Management.

4.9 Quality Assurance/Quality Control

All data entered into NPSTORET should be reviewed for accuracy by the individual entering the data. The project manager is responsible for ensuring all QA/QC procedures are followed, including proper data entry, validation, and documentation. See SOP #12 for more details on QA/QC for data entry and management.

4.11 Literature Cited

- Hart, M., and U. Gafvert (editors). 2005. Data management plan: Great Lakes Inventory and Monitoring Network. National Park Service Great Lakes Inventory and Monitoring Network Report GLKN/2005/20. National Park Service, Ashland, Wisconsin.
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Standard Operating Procedure #5: Decontamination of Equipment to Remove Exotic Species

Version 1.2

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Revision History Log

The following table lists all edits and amendments to this document since the original publication date. Information entered in the log must be complete and concise. Users of this standard operating procedure will promptly notify the project manager and/or the Great Lakes Network (GLKN) data manager about recommended and required changes. The project manager must review and incorporate all changes, complete the revision history log, and change the date and version number on the title page and in the header of the document file. For complete instructions, please refer to Revising the Protocol, SOP #13.

Previous Version Number*	Revision Date	Author (with title and affiliation)	Location in Document and Concise Description of Revision	Reason for Change	New Version Number
1.0	3/1/2010	Richard Damstra, Joan Elias GLKN Aquatic Ecologists	Minor clarifications throughout; section 5.2.2, Procedures for Decontamination updated such that a single procedure is outlined regardless of remoteness of site; Appendix B, HACCP updated to include zebra mussels expansion	To clarify protocol and bring field procedures and protocol into agreement, based on staff field experience.	1.1
1.1	5/1/2015	David VanderMeulen, Rick Damstra GLKN	Updated the NPS contact author and authorship, fixed a few grammatical errors; Appendix B, HACCP updated to reflect changes in sites/parks sampled, included Asian carp, and Eurasian water milfoil expansion.	To update contact author since Version 1.1 was published in 2010; To clarify protocol and bring field procedures and protocol into agreement, based on staff field experience.	1.2

*Add rows as needed for each change or set of changes tied to an updated version number

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I wish to thank Jeff Gunderson and Doug Jensen, of the Minnesota Sea Grant Program, Phil Moy, of the Wisconsin Sea Grant Program, and Mike Klepinger, of the Michigan Sea Grant Program, for their willingness to discuss exotic species issues in national parks and for providing information on the Hazard Analysis Critical Control Point (HACCP) method.

5.0 Introduction

Aquatic invasive species (AIS) are an issue of increasing concern nationwide. In the Great Lakes, for example, approximately 140 exotic species have invaded since the late 1800s (Great Lakes Information Network 2005) and have had enormous ecosystem effects (e.g., Dermott and Kerec 1997, Barnhisel and Kerfoot 2004, Hoff 2004, Great Lakes Commission 2005). Some of these species have invaded inland water bodies, where disruptions to native species are also documented (e.g., Jack and Thorp 2000, Indrissi et al. 2001, Compton and Kerfoot 2004). Zebra mussels (*Dreissena polymorpha*) now occur in several lakes at Sleeping Bear Dunes National Lakeshore, and the spiny waterflea (*Bythotrephes cederstroemi*) has been found in Grand Sable Lake, Pictured Rocks National Lakeshore. Invasive plants associated with waterbodies, such as hybrid cattails (*Typha glauca* x *Typha angustifolia*), are found at many Great Lakes Network parks, including Isle Royale. Bighead carp (*Hypophthalmichthys nobliis*) were caught or captured in Lake St. Croix at the St. Croix National Scenic Riverway and in the Minnesota River, a large tributary to the Mississippi River within the boundaries of Mississippi National River and Recreation Area. When Network and park staff conduct routine water quality monitoring, it is important to ensure these aquatic exotics or other emerging threats are not transferred from contaminated water bodies to uncontaminated water bodies.

As we conduct water quality monitoring on large rivers, we are concerned with the limited suite of AIS that can adhere to, or passively catch on, sampling equipment (including waders, boats and trailers). In this standard operating procedure (SOP) we will focus on aquatic plants, zooplankton, and mussels.

At each park, we will use the Hazard Analysis Critical Control Point (HACCP) method (Gunderson and Kinuunen 2004; Appendix A) to identify water bodies at risk and locations where control of AIS is imperative. It is critical to integrate this SOP with monitoring for the occurrence of AIS in large rivers.

5.1 Equipment and Supplies

Sampling equipment used in monitoring large rivers is detailed in SOP #6. We designed the decontamination process to be simple, straight-forward, and consistent with NPS wilderness considerations. Specifically, no toxic chemicals are required for decontamination.

The supplies and equipment required for decontamination of sampling equipment between sites consists of:

- tap water
- hose and sprayer nozzle
- large pot
- portable propane burner (sold as a turkey deep fryer) or other means of heating water in a large pot (Figure 5-1)

- thermometer for measuring temperature of heated water
- scrub brushes



Figure 5-1. An example of a propane burner and pot used to heat water to 140° F (60° C) for decontaminating non-electronic equipment. Any large pot and heat source is acceptable; however a thermometer should be used to verify that sufficient temperatures have been achieved.

5.2 Decontamination Procedures

5.2.1 Aquatic Invasive Species Procedure Plan and Hazard Analysis Worksheet

The aquatic invasive species hazard analysis and critical control point (HACCP) worksheet is used in describing the sampling project, including types of gear, methods of transportation, steps involved in the procedure; and analyzing the potential risk of AIS transport from one water body to another, or one section of a stream to another section. Details on completing the worksheet are described in the AIS-HACCP manual (Gunderson and Kinuunen 2004), and are included in Appendix A. The example worksheet included in Appendix B is for St. Croix National Scenic Riverway. The Network has modified the worksheet slightly to suit our needs. Worksheets will be completed for each park unit, and will be updated annually prior to the sampling season. Worksheets will also be updated in the following circumstances:

- change in sampling equipment or techniques

- addition of a new water body or to the sampling regime
- change in the order of sampling water bodies
- detection of a new AIS in a water body
- change in scientific knowledge regarding life history, potential hazard, or control of an AIS.

5.2.2 Procedures for Decontamination

Many AIS, especially plants, can be controlled by simply being vigilant and periodically examining equipment, clothing, and boots and removing materials suspected to be an AIS species (cattail “fluff” or other seeds, viable plant materials, animal “hitchhikers” such as spiny water flea stuck or other invertebrates stuck on equipment).

When the hazard analysis worksheet identifies a significant AIS risk and adequate drying or freezing of equipment is not feasible (five days in the sun for boats and trailers, 10 days for other equipment, or two days of freezing; Gunderson and Kinuunen 2004), the procedures below will be followed. The decontamination procedure assumes tap water is available.

Basic guidelines:

- Be vigilant about periodically checking equipment, clothing, and boots during portages and especially upon arrival and departure from sampling sites. Look for signs of AIS, especially plant materials.
- When possible, avoid using equipment on both infested and uninfested waters. Designate a set of equipment for both types of waters.
- Always begin sampling with sites not known to harbor exotic species; sample sites known or suspected to contain exotic species last.
- Thoroughly clean or disinfect all equipment between uses in all inland waters.
- When possible, allow sampling equipment, to dry for 10 days after sampling a water body known to be infested with AIS.
- Due to emerging science on the transport of AIS in felt soles of wading boots, lug soles constructed of a non-porous material, such as rubber, should be utilized for ease of decontamination (Gates 2007, Bothwell et al. 2009).

Step-by-Step Procedure:

When finished at a sampling site and before sampling the next site:

- Remove or rinse mud, plant material, invertebrates, and other visible organic material from boots, nets, and all sampling equipment with a hose and high pressure nozzle.
- Non-electronic gear, such as lines, tapes, dipnets, wading boots, should be exposed to water heated to 140°F (60°C) and held at this temperature for at least 1 minute. Portable turkey fryers are ideal for this, although a large pot on a stove will work. Rags and brushes used in decontamination procedures should be heat treated as well. Electronic equipment such as

flow meters, depth probes, temperature probes, data collection sondes, and associated cables should be rinsed with copious amounts of tap water and cleaned with brushes and/or clean rags.

- Pay particular attention to lines, cracks, and crevices in equipment, and rinse well.
- Visually inspect all gear to verify that all equipment and gear has been cleaned and rinsed.
- Take measures to allow equipment to dry out as completely as possible.

5.3 Documentation

The original AIS-HACCP worksheet and all updates will be maintained at each park and copies will be kept at the Network office. Documents used to complete each worksheet will also be kept on file at the parks and at the Network. Examples of such supporting documentation include information used in analyzing the hazards of contamination and determining adequate prevention of AIS spread, and the most current geographic range of AIS.

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Appendix 5A. Instructions for completing a Hazard Analysis and Critical Control Point Plan.

The following instructions for completing HACCP Procedure Form, Potential Hazards Worksheet, Hazard Analysis Worksheet, and Plan are taken directly from the AIS-HACCP manual (Gunderson and Kinuunen 2004). Not all steps will apply to the Great Lakes Inventory and Monitoring Network's protocol for decontamination of water quality monitoring field equipment and gear.

Preliminary Steps

1: Document general information.

Record the name and address of your facility or agency in the spaces provided on the first page of the Hazard Analysis Worksheet and the AISHACCP Plan Form (Appendix 9).

2: Describe the cultured or wild harvested fish (if applicable).

Identify the market name or Latin name (species) of the fish. Examples:

- Fathead minnows (*Pimephales promelas*)
- Golden shiners (*Notemigonus crysoleucas*)
- White sucker (*Catostomus commersoni*)
- Walleye (*Sander vitreus*)

Fully describe the product. Examples:

- Fathead minnows graded on 16 grader
- Golden shiners graded on 21 grader
- White suckers ungraded
- Rosy red minnows from Arkansas, held in ponds until distribution
- White suckers graded on a 23 grader
- Walleye fingerlings 5–8 inches
- Yellow perch fingerlings 2.5–3 inches

For management, research, and enforcement activities:

- Completely describe research or management activities
- Describe all equipment and gear that will be used
- Identify when and how activities will be conducted

Record this information in the spaces provided on the first page of the Hazard Analysis Worksheet and the AIS-HACCP Plan Form.

3: Describe the harvest, production, management, research, or enforcement activity.

Identify how the product or the samples are collected, stored, and distributed. Identify whether any special shipping or handling methods are used. Examples:

- Wild harvested with seines, held in tanks, graded, then trucked to retail stores
- Pond-raised, seined, then held in different ponds over the winter, trucked to retail stores
- Pond-raised, trapped from ponds, then transferred directly to lakes for stocking
- Anglers on three lakes are checked for violations each day for a given period of time
- Electroshocking and seining are conducted on 10 lakes to assess year class strength

Record this information in the spaces provided on the first page of the Hazard Analysis Worksheet and the AIS-HACCP Plan Form.

4: Identify the intended use and customer (if applicable).

Identify how the product will be used. Examples:

- Live fishing bait
- Feeder fish (feeding pond or aquarium fish)
- Stocking into public waters
- Stocking into private waters
- Stocking into aquaculture production facility (indoor or outdoor)
- Scale and stomach samples will be brought back to the office for analysis
- Fish will be brought back for captive brood stock

Identify your intended customer or user of the product. Examples:

- General public
- A wholesaler
- A retail store
- Fish farmer
- State agency

Record this information in the spaces provided on the first page of the Hazard Analysis Worksheet and the AIS-HACCP Plan Form.

5: Develop a flow diagram.

The purpose of the flow diagram is to provide a clear, simple description of the steps involved in producing your fishery products or conducting your management, research, and enforcement activities. The flow diagram should cover all of the steps in the process that your firm or agency performs. The flow diagram should be verified on-site for accuracy. Examples of flow diagrams can be found in appendices 3–8.

Hazard Analysis Worksheet

6: Set up the Hazard Analysis Worksheet.

Record each of the steps from the flow diagram in column 1 of the Hazard Analysis Worksheet.

7: Identify the potential AIS-related hazards.

Record the AIS-related hazards for each step. Use your own expertise and that of others to identify AIS hazards related to your fish production, or management, research, or enforcement practices. Check with appropriate agencies to determine if the waters in which you conduct activities are infested with invasive species.

Even if you think you have effective hazard controls in place, record the hazard. For example, your equipment might be free of AIS plant material because of the: 1) absence of AIS in the area of harvest or sampling in an infested water body; or 2) existence of inadvertent hazard controls (procedures you typically use in the course of your activities that may remove AIS).

8: Complete the Hazard Analysis Worksheet.

Completing the Hazard Analysis Worksheet requires understanding potential hazards, determining if each potential hazard is significant, and identifying critical control points for each significant hazard associated with your product or activities.

AIS –HACCP Plan Form

9: Complete the AIS-HACCP Plan Form.

Copy the Critical Control Points from column 6 of the Hazard Analysis Worksheet to column 1 of the AIS-HACCP Plan Form. Enter the associated hazard(s) from column 2 of the worksheet to column 2 of the plan form. If you did not identify significant hazards and CCPs, you do not need to complete an AIS-HACCP plan.

Complete the AIS-HACCP Plan Form by designing techniques, methods, and treatments to deal with each significant hazard in column 2. For each significant hazard:

- set critical limits
- establish monitoring procedures
- establish corrective action procedures
- establish a record keeping system
- establish verification procedures

After you completed these steps for each hazard, the AIS-HACCP plan form is finished.

To signify that the AIS-HACCP plan has been accepted for implementation, the responsible individual on-site or a higher level official should sign and date the first page of the plan form.

Appendix 5B. Aquatic Invasive Species Hazard Analysis and Critical Control Point Plan for water quality monitoring of Large Rivers at St. Croix National Scenic Riverway (version 1.2).

AIS-HACCP PLAN

Aquatic Invasive Species-Hazard Analysis and Critical Control Point

Procedure Form

Procedure Description

Park info:	<i>Park name: St. Croix National Scenic River</i>			
	<i>Address: 401 N Hamilton St</i>	<i>City: St. Croix Falls</i>	<i>State: WI</i>	<i>Zip: 54024</i>
(if applicable):	<i>Fish species: NA</i>			
Management, research, or monitoring activity:	<i>Activity: Water Quality Monitoring</i>			
Method of transportation, distribution, and storage of boats, gear, etc.	<i>Methods: Boat and gear either trailered or portaged to public landing sites. Profiles of water temperature, dissolved oxygen, specific conductance, and pH are conducted with multi-probe sonde at all sites. Water is collected for lab analysis by hand with a jug, or with an integrated tube or Van Dorn sampler. Multiple locations on the river or its tributaries may be sampled in one day.</i>			
Procedure Flow	<i>(1)Day 1: NPS staff drives to Phipps Landing. Conduct water quality sampling by wading into the Namekagon River. Collect a grab sample of water, collect water quality data with multiprobe and through field observations. Travel to Earl Landing.</i>			
List the steps involved in the management, research, or monitoring activity.	<i>(2)Day 1 continued: Conduct water quality sampling on the Namekagon River by working off the bridge on North Rd., just upstream from the Earl Landing. Collect a grab sample of water using a Van Dorn sampler or equivalent piece of gear, collect water quality data with multiprobe from bridge and through field observations. Travel to Namekagon River at Namekagon Trail Bridge.</i>			
	<i>(3)Day 1 continued: Conduct water quality monitoring on the Namekagon River by working off the Namekagon Trail Bridge. Collect a grab sample of water using a Van Dorn sampler or equivalent piece of gear, collect water quality data with a mutilprobe from bridge and through field observations. Travel to CCC Bridge.</i>			
	<i>(4)Day 1 continued: Conduct water quality monitoring on the Upper St. Croix River by working off the CCC Bridge. Collect a grab sample of water using a Van Dorn sampler or equivalent peice of gear, collect water quality data with a mutilprobe from bridge and through field observations. Travel to lab and process samples.</i>			

(continued on back)

(5)Day 2: NPS staff drives to Norway Point Landing on the St. Croix River towing a boat and trailer. Launch boat, travel upstream to sample site, conduct water quality monitoring from the boat. Collect a water sample via a grab sample or integrating tube, depending on water depth. Collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to Highway 70 boat launch.

(6)Day 2 continued: NPS staff drives to Highway 70 Landing on the St. Croix River towing a boat and trailer. Launch boat, travel to midstream to sample site, conduct water quality monitoring from the boat. Collect a water sample via a grab sample or integrating tube, depending on water depth. Collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to the Nevers Dam boat launch.

(7)Day 2 continued: Conduct water quality monitoring on the St. Croix River by launching a boat at the Nevers Dam boat launch. Boat upstream to sample site, conduct water quality sampling from boat. Collect a water sample via grab sample or by using integrating tube, depending on water depth. Collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to lab and process samples.

(8)Day 3: Conduct water quality monitoring on Lake St. Croix near Bayport, WI by launching boat at the public boat launch at Bayport. Boat to sample site, conduct water quality sampling from boat. Collect an integrated sample using an integrating tube, collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to Hudson, WI boat launch.

(9)Day 3 continued: Conduct water quality sampling on the St. Croix River by launching boat at the public boat launch in Hudson, WI. Boat to sample site, conduct water quality sampling from boat. Collect an integrated sample using an integrating tube, collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to Prescott, WI boat launch.

(10)Day 3 continued: Conduct water quality monitoring on the Lake St. Croix near Prescott, WI by launching boat at the public boat launch on the Mississippi River in Prescott, WI. Travel up the St. Croix River to sample site, conduct water quality sampling from boat. Collect an integrated sample using an integrating tube, collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel back to lab and process samples.

Upon
completion of
the AIS-
HACCP plan,
sign here to
accept plan for
implementation.

Name: _____

Signature: _____

Date: _____

AIS-HACCP PLAN

Aquatic Invasive Species-Hazard Analysis and Critical Control Point

Potential Hazards Worksheet

Potential AIS Hazards

List all relevant species

AIS Fish and Other Vertebrates Examples: round goby, ruffe, carp, etc.	<i>Bighead carp (Hypophthalmichthys nobilis) captured in Lake St. Croix as far upstream as Bayport, MN.</i>
AIS Invertebrates Examples: zebra mussel, spiny water flea, rusty crayfish, etc.	<i>From the Stillwater lift bridge to the confluence with the Mississippi River is declared infested by zebra mussels (Dreissena polymorpha) by the MNDNR and WIDNR. Rusty crayfish (Orconectes rusticus) have also been found from Fox Landing to the confluence and in the Snake, Apple, and Kinnickinnic Rivers.</i>
AIS Plants Examples: curly pondweed, Eurasian water-milfoil, etc.	<i>The St. Croix River in Washington County is designated as infested by Eurasian watermilfoil (Myriophyllum spicatum) by the MNDNR. Gordon Dam Flowage on the Upper St. Croix was treated for Eurasian water millfoil in 2015.</i>
AIS Pathogens Examples: whirling disease, <i>Heterosporis</i>, etc.	N/A

Next Step:

After identifying the potential hazards, complete a hazard analysis form.

AIS-HACCP PLAN**Aquatic Invasive Species-Hazard Analysis and Critical Control Point****Hazard Analysis Worksheet**

(1) Activity	(2) Hazards	(3) Significance	(4) Justification	(5) Control	(6) CCP
Activity step from flow diagram, page 1	Potential AIS hazards introduced or controlled at this step (from potential hazards worksheet)	Are AIS hazards significant? (Yes/No)	Justify your decisions for column 3.	What control measures will be applied to prevent the significant hazards?	Is this step a critical control point? (Yes/No)
<i>Work Flow Step (1) Day 1: NPS staff drives to Phipps Landing. Conduct water quality sampling by wading into the Namekagon River. Collect a grab sample of water, collect water quality data with multiprobe and through field observations. Travel to Earl Landing.</i>	Fish/Other Vertebrates N/A	No	Equipment used will not capture any fish.		No
	Invertebrate N/A	No	No invertebrate AIS on this stretch of the river.		No
	Plant N/A	No	No plant AIS at site.		No
	Pathogens N/A	No	AIS pathogens not present.		No

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AIS-HACCP PLAN**Aquatic Invasive Species-Hazard Analysis and Critical Control Point****Hazard Analysis Worksheet**

(1) Activity	(2) Hazards	(3) Significance	(4) Justification	(5) Control	(6) CCP
Activity step from flow diagram, page 1	Potential AIS hazards introduced or controlled at this step (from potential hazards worksheet)	Are AIS hazards significant? (Yes/No)	Justify your decisions for column 3.	What control measures will be applied to prevent the significant hazards?	Is this step a critical control point? (Yes/No)
(2)Day 1 continued: Conduct water quality sampling on the Namekagon River by working off the bridge on North Rd., just upstream from the Earl Landing. Collect a grab sample of water using a Van Dorn sampler or equivalent piece of gear, collect water quality data with multiprobe from bridge and through field observations. Travel to Namekagon River at Namekagon Trail Bridge.	Fish/Other Vertebrates N/A	No	Equipment used will not capture any fish.		No
	Invertebrate N/A	No	No invertebrate AIS on this stretch of the river.		No
	Plant N/A	No	No plant AIS at site.		No
	Pathogens N/A	No	AIS pathogens not present		No

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AIS-HACCP PLAN***Aquatic Invasive Species-Hazard Analysis and Critical Control Point******Hazard Analysis Worksheet***

(1)	(2)	(3)	(4)	(5)	(6)
Activity	Hazards	Significance	Justification	Control	CCP
Activity step from flow diagram, page 1	Potential AIS hazards introduced or controlled at this step (from potential hazards worksheet)	Are AIS hazards significant? (Yes/No)	Justify your decisions for column 3.	What control measures will be applied to prevent the significant hazards?	Is this step a critical control point? (Yes/No)
(3)Day 1 continued: Conduct water quality monitoring on the Namekagon River by working off the Namekagon Trail Bridge. Collect a grab sample of water using a Van Dorn sampler or equivalent piece of gear, collect water quality data with a mutilprobe from bridge and through field observations. Travel to CCC Bridge.	Fish/Other Vertebrates N/A	No	Boat and equipment used will not capture any fish.		No
	Invertebrate N/A	No	No invertebrate AIS on this stretch of the river		No
	Plant N/A	No	No plant AIS at site.		No
	Pathogens N/A	No	AIS pathogens not present.		No

SOP 5-19

AIS-HACCP PLAN***Aquatic Invasive Species-Hazard Analysis and Critical Control Point******Hazard Analysis Worksheet***

(1)	(2)	(3)	(4)	(5)	(6)
Activity	Hazards	Significance	Justification	Control	CCP
Activity step from flow diagram, page 1	Potential AIS hazards introduced or controlled at this step (from potential hazards worksheet)	Are AIS hazards significant? (Yes/No)	Justify your decisions for column 3.	What control measures will be applied to prevent the significant hazards?	Is this step a critical control point? (Yes/No)
(4)Day 1 continued: Conduct water quality monitoring on the Upper St. Croix River by working off the CCC Bridge. Collect a grab sample of water using a Van Dorn sampler or equivalent piece of gear, collect water quality data with a mutilprobe from bridge and through field observations. Travel to lab and process samples.	Fish/Other Vertebrates N/A	No	Boat and equipment used will not capture any fish.		No
	Invertebrate Rusty Crayfish	No	No invertebrate AIS on this stretch of the river		No
	Plant Eurasian water millfoil	Yes	No plant AIS at site, but equipment will be checked since Eurasian water millfoil is found upstream from site at Gordon Dam Flowage.		Yes
	Pathogens N/A	No	AIS pathogens not present.		No

SOP 5-20

AIS-HACCP PLAN**Aquatic Invasive Species-Hazard Analysis and Critical Control Point****Hazard Analysis Worksheet**

(1)	(2)	(3)	(4)	(5)	(6)
Activity	Hazards	Significance	Justification	Control	CCP
Activity step from flow diagram, page 1	Potential AIS hazards introduced or controlled at this step (from potential hazards worksheet)	Are AIS hazards significant? (Yes/No)	Justify your decisions for column 3.	What control measures will be applied to prevent the significant hazards?	Is this step a critical control point? (Yes/No)
(5)Day 2: NPS staff drives to Norway Point Landing on the St. Croix River towing a boat and trailer. Launch boat, travel upstream to sample site, conduct water quality monitoring from the boat. Collect a water sample via a grab sample or integrating tube, depending on water depth. Collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to Highway 70 boat launch.	Fish/Other Vertebrates N/A	No	Boat and equipment used will not capture any fish.		No
	Invertebrate Rusty Crayfish	No	No invertebrate AIS on this stretch of the river		No
	Plant Eurasian watermilfoil	No	No plant AIS at site.		No
	Pathogens N/A	No	AIS pathogens not present.		No

SOP 5-21

AIS-HACCP PLAN**Aquatic Invasive Species-Hazard Analysis and Critical Control Point****Hazard Analysis Worksheet**

(1)	(2)	(3)	(4)	(5)	(6)
Activity	Hazards	Significance	Justification	Control	CCP
Activity step from flow diagram, page 1	Potential AIS hazards introduced or controlled at this step (from potential hazards worksheet)	Are AIS hazards significant? (Yes/No)	Justify your decisions for column 3.	What control measures will be applied to prevent the significant hazards?	Is this step a critical control point? (Yes/No)
(6)Day 2 continued: NPS staff drives to Highway 70 Landing on the St. Croix River towing a boat and trailer. Launch boat, travel to midstream to sample site, conduct water quality monitoring from the boat. Collect a water sample via a grab sample or integrating tube, depending on water depth. Collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to the Nevers Dam boat launch.	Fish/Other Vertebrates N/A	No	Boat and equipment used will not capture any fish.		No
	Invertebrate Rusty Crayfish	Yes	Rusty crayfish present.	Inspect boat for stowaway rusty crayfish.	Yes
	Plant Eurasian watermilfoil	No	No plant AIS at site.		No
	Pathogens N/A	No	AIS pathogens not present.		No

SOP 5-22

AIS-HACCP PLAN**Aquatic Invasive Species-Hazard Analysis and Critical Control Point****Hazard Analysis Worksheet**

(1)	(2)	(3)	(4)	(5)	(6)
Activity	Hazards	Significance	Justification	Control	CCP
Activity step from flow diagram, page 1	Potential AIS hazards introduced or controlled at this step (from potential hazards worksheet)	Are AIS hazards significant? (Yes/No)	Justify your decisions for column 3.	What control measures will be applied to prevent the significant hazards?	Is this step a critical control point? (Yes/No)
(7)Day 2 continued: Conduct water quality monitoring on the St. Croix River by launching a boat at the Nevers Dam boat launch. Boat upstream to sample site, conduct water quality sampling from boat. Collect a water sample via grab sample or by using integrating tube, depending on water depth. Collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to lab and process samples.	Fish/Other Vertebrates N/A	No	Boat and equipment used will not capture any fish.		No
	Invertebrate Rusty Crayfish	Yes	Rusty crayfish present.	Inspect boat for stowaway rusty crayfish.	Yes
	Plant Eurasian watermilfoil	No	No plant AIS at site.		No
	Pathogens N/A	No	AIS pathogens not present.		No

SOP 5-23

AIS-HACCP PLAN**Aquatic Invasive Species-Hazard Analysis and Critical Control Point****Hazard Analysis Worksheet**

(1) Activity	(2) Hazards	(3) Significance	(4) Justification	(5) Control	(6) CCP
Activity step from flow diagram, page 1	Potential AIS hazards introduced or controlled at this step (from potential hazards worksheet)	Are AIS hazards significant? (Yes/No)	Justify your decisions for column 3.	What control measures will be applied to prevent the significant hazards?	Is this step a critical control point? (Yes/No)
<i>(8)Day 3: Conduct water quality monitoring on Lake St. Croix near Bayport, WI by launching boat at the public boat launch at Bayport. Boat to sample site, conduct water quality sampling from boat. Collect an integrated sample using an integrating tube, collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to Hudson, WI boat launch.</i>	Fish/Other Vertebrates Asian carp	No	Boat and equipment used will not capture any fish.		No
	Invertebrates Zebra Mussels, Rusty Crayfish	Yes	Zebra mussel veligers may attach to equipment and boat. Rusty crayfish present.	If traveling to non-infested area, boat and equipment should be sprayed with hot water (140F); Inspect boat for stowaway rusty crayfish.	Yes
	Plant Eurasian Watermilfoil	Yes	Eurasian watermilfoil infested waters.	Inspect boat, anchor, and rope before and after sampling; remove any mud and plant material	Yes
	Pathogens N/A	No	AIS pathogens not present.		No

SOP 5-24

AIS-HACCP PLAN**Aquatic Invasive Species-Hazard Analysis and Critical Control Point****Hazard Analysis Worksheet**

(1) Activity	(2) Hazards	(3) Significance	(4) Justification	(5) Control	(6) CCP
Activity step from flow diagram, page 1	Potential AIS hazards introduced or controlled at this step (from potential hazards worksheet)	Are AIS hazards significant? (Yes/No)	Justify your decisions for column 3.	What control measures will be applied to prevent the significant hazards?	Is this step a critical control point? (Yes/No)
(9)Day 3 continued: Conduct water quality sampling on the St. Croix River by launching boat at the public boat launch in Hudson, WI. Boat to sample site, conduct water quality sampling from boat. Collect an integrated sample using an integrating tube, collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to Prescott, WI boat launch.	Fish/Other Vertebrates Asian carp	No	Boat and equipment used will not capture any fish.		No
	Invertebrate Zebra Mussels, Rusty Crayfish	Yes	Zebra mussel veligers may attach to equipment and boat. Rusty crayfish present.	If traveling to non-infested area, boat and equipment should be sprayed with hot water (140F); Inspect boat for stowaway rusty crayfish.	Yes
	Plant Eurasian Watermilfoil	Yes	Eurasian watermilfoil infested waters.	Inspect boat, anchor, and rope before and after sampling; remove any mud and plant material	Yes
	Pathogens N/A	No	AIS pathogens not present.		No

SOP 5-25

AIS-HACCP PLAN**Aquatic Invasive Species-Hazard Analysis and Critical Control Point****Hazard Analysis Worksheet**

(1)	(2)	(3)	(4)	(5)	(6)
Activity	Hazards	Significance	Justification	Control	CCP
Activity step from flow diagram, page 1	Potential AIS hazards introduced or controlled at this step (from potential hazards worksheet)	Are AIS hazards significant? (Yes/No)	Justify your decisions for column 3.	What control measures will be applied to prevent the significant hazards?	Is this step a critical control point? (Yes/No)
(10)Day 3 continued: Conduct water quality monitoring on the Lake St. Croix near Prescott, WI by launching boat at the public boat launch on the Mississippi River in Prescott, WI. Travel up the St. Croix River to sample site, conduct water quality sampling from boat. Collect an integrated sample using an integrating tube, collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel back to lab and process samples.	Fish/Other Vertebrates Asian carp	No	Boat and equipment used will not capture any fish.		No
	Invertebrate Zebra Mussels, Rusty Crayfish	Yes	Zebra mussel veligers may attach to equipment and boat, Rusty crayfish present.	If traveling to non-infested area, boat and equipment should be sprayed with hot water (140F); Inspect boat for stowaway rusty crayfish.	Yes
	Plant Eurasian Watermilfoil	Yes	Eurasian watermilfoil infested waters.	Inspect boat, anchor, and rope before and after sampling; remove any mud and plant material.	Yes
	Pathogens N/A	No	AIS pathogens not present.		No

SOP 5-26

AIS-HACCP PLAN

Aquatic Invasive Species-Hazard Analysis and Critical Control Point

AIS-HACCP Plan Form

Critical Control Point Each row answered "yes" in column 6 of Hazard Analysis Form	(4)Day 1 continued: Conduct water quality monitoring on the Upper St. Croix River by working off the CCC Bridge. Collect a grab sample of water using a Van Dorn sampler or equivalent piece of gear, collect water quality data with a multiprobe from bridge and through field observations. Travel to lab and process samples.	(6)Day 2 continued: NPS staff drives to Highway 70 Landing on the St. Croix River towing a boat and trailer. Launch boat, travel to midstream to sample site, conduct water quality monitoring from the boat. Collect a water sample via a grab sample or integrating tube, depending on water depth. Collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to the Nevers Dam boat launch.
Significant Hazards as determined in column 3 of Hazard Analysis Form	(2) The Gordon Dam Flowage upstream from this site is infested with Eurasian water millfoil.	(2) Rusty crayfish present.
Limits for Each Control Measure	(3) No viable plant parts remain on boat, trailer, equipment, or waders.	(3) Rusty crayfish remain on boat, trailer, equipment, or waders.
Monitoring Describe what is being monitored	(4) Presence of plant material. Ensure that boat, trailer, waders, and all equipment are free of viable plant parts.	(4) Rusty crayfish. Ensure the boat, trailer, waders, and other equipment are free of rusty crayfish.
Explain how the monitoring will take place	(5) Visual inspection of boat, trailer, waders, and other equipment for plant material.	(5) Visual inspection of boat, trailer, waders, and other equipment for rusty crayfish.
Frequency of monitoring	(6) Each time, before and after boat, trailer, and equipment is used at site.	(6) Each time, before and after boat, trailer, and equipment is used at site.
Person or position responsible for monitoring	(7) Water quality monitoring staff.	(7) Water quality monitoring staff.
Corrective Actions Actions taken when limits of control measures are not met	(8) Cease operation and clean boat, trailer, waders, and equipment before launching onto water.	(8) Cease operation and clean boat, trailer, waders, and equipment before launching onto water.
Verification Method of verification	(9) Review records.	(9) Review records.
Records List what is recorded at each critical control point	(10) Record the procedure used to remove plant fragments. Record that inspections occurred prior to leaving one site and entering the next site.	(10) Record the procedure used to remove and rusty crayfish. Record that inspections occurred prior to leaving one launch and before entering the river at the next site

AIS-HACCP PLAN

Aquatic Invasive Species-Hazard Analysis and Critical Control Point AIS-HACCP Plan Form (continued)

Critical Control Point Each row answered "yes" in column 6 of Hazard Analysis Form	<i>(7) Day 2 continued: Conduct water quality monitoring on the St. Croix River by launching a boat at the Nevers Dam boat launch. Boat upstream to sample site, conduct water quality sampling from boat. Collect a water sample via grab sample or by using integrating tube, depending on water depth. Collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to lab and process samples.</i>	<i>Work Flow Step (10): Conduct water quality monitoring on Lake St. Croix near Bayport, WI by launching boat at the public boat launch at Bayport. Boat to sample site, conduct water quality sampling from boat. Collect an integrated sample using an integrating tube, collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel back to lab, process water samples.</i>
Significant Hazards as determined in column 3 of Hazard Analysis Form	<i>(2) Rusty crayfish present.</i>	<i>(2) Zebra mussels and rusty crayfish present.</i>
Limits for Each Control Measure	<i>(3) Rusty crayfish remain on boat, trailer, equipment, or waders.</i>	<i>(3) No zebra mussels or rusty crayfish remain on boat, trailer, equipment, or waders.</i>
Monitoring Describe what is being monitored	<i>(4) Rusty crayfish. Ensure the boat, trailer, waders, and other equipment are free of rusty crayfish.</i>	<i>(4) Presence of zebra mussels and rusty crayfish. Ensure the boat, trailer, waders, and other equipment are free of rusty crayfish and zebra mussels.</i>
Explain how the monitoring will take place	<i>(5) Visual inspection of boat, trailer, waders, and other equipment for rusty crayfish.</i>	<i>(5) Visual inspection of boat, trailer, waders, and other equipment for rusty crayfish and zebra mussels.</i>
Frequency of monitoring	<i>(6) Each time, before and after boat, trailer, and equipment is used at site.</i>	<i>(6) Each time, before and after boat, trailer, and equipment is used at site.</i>
Person or position responsible for monitoring	<i>(7) Water quality monitoring staff.</i>	<i>(7) Water quality monitoring staff.</i>
Corrective Actions Actions taken when limits of control measures are not met	(8) Cease operation and clean boat, trailer, waders, and equipment before launching onto water.	(8) Cease operation and clean boat, trailer, waders, and equipment before launching onto water.
Verification Method of verification	<i>(9) Review records.</i>	<i>(9) Review records.</i>
Records List what is recorded at each critical control point	<i>(10) Record the procedure used to remove and rusty crayfish. Record that inspections occurred prior to leaving one launch and before entering the river at the next site.</i>	<i>(10) Record the procedure used to remove zebra mussels and rusty crayfish. Record that inspections occurred prior to leaving one launch and before entering the river at the next site.</i>

This form accommodates two Critical Control points. Attach additional pages of this form as necessary.

Final Step:

Once the HACCP plan is completed, attach it to the signed procedure form with the hazard analysis worksheets.

AIS-HACCP PLAN

Aquatic Invasive Species-Hazard Analysis and Critical Control Point AIS-HACCP Plan Form (continued)

Critical Control Point Each row answered "yes" in column 6 of Hazard Analysis Form	<i>Work Flow Step (10): Conduct water quality monitoring on Lake St. Croix near Bayport, WI by launching boat at the public boat launch at Bayport. Boat to sample site, conduct water quality sampling from boat. Collect an integrated sample using an integrating tube, collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel back to lab, process water samples.</i>	<i>(9)Day 3 continued: Conduct water quality sampling on the St. Croix River by launching boat at the public boat launch in Hudson, WI. Boat to sample site, conduct water quality sampling from boat. Collect an integrated sample using an integrating tube, collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to Prescott, WI boat launch.</i>
Significant Hazards as determined in column 3 of Hazard Analysis Form	(2) The St. Croix River in Washington County has been designated as infested with Eurasian watermilfoil by the MNDNR.	(2) Zebra mussels and rusty crayfish present
Limits for Each Control Measure	(3) No viable plant parts remain on boat, trailer, equipment, or waders.	(3) No zebra mussels or rusty crayfish remain on boat, trailer, equipment, or waders.
Monitoring Describe what is being monitored	(4) Presence of plant material. Ensure that boat, trailer, waders, and all equipment are free of viable plant parts.	(4) Presence of zebra mussels and rusty crayfish. Ensure the boat, trailer, waders, and other equipment are free of rusty crayfish and zebra mussels.
Explain how the monitoring will take place	(5) Visual inspection of boat, trailer, waders, and other equipment for plant material.	(5) Visual inspection of boat, trailer, waders, and other equipment for rusty crayfish and zebra mussels.
Frequency of monitoring	(6) Each time, before and after boat, trailer, and equipment is used at site.	(6) Each time, before and after boat, trailer, and equipment is used at site.
Person or position responsible for monitoring	(7) Water quality monitoring staff.	(7) Water quality monitoring staff.
Corrective Actions Actions taken when limits of control measures are not met	(8) Cease operation and clean boat, trailer, waders, and equipment before launching onto water.	(8) Cease operation and clean boat, trailer, waders, and equipment before launching onto water.
Verification Method of verification	(9) Review records.	(9) Review records.
Records List what is recorded at each critical control point	(10) Record the procedure used to remove plant fragments. Record that inspections occurred prior to leaving one site and entering the next site.	(10) Record the procedure used to remove zebra mussels and rusty crayfish. Record that inspections occurred prior to leaving one launch and before entering the river at the next site.

This form accommodates two Critical Control points. Attach additional pages of this form as necessary.

Final Step:

Once the HACCP plan is completed, attach it to the signed procedure form with the hazard analysis worksheets.

AIS-HACCP PLAN

Aquatic Invasive Species-Hazard Analysis and Critical Control Point AIS-HACCP Plan Form (continued)

Critical Control Point Each row answered "yes" in column 6 of Hazard Analysis Form	(9)Day 3 continued: Conduct water quality sampling on the St. Croix River by launching boat at the public boat launch in Hudson, WI. Boat to sample site, conduct water quality sampling from boat. Collect an integrated sample using an integrating tube, collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel to Prescott, WI boat launch.	(10)Day 3 continued: Conduct water quality monitoring on the Lake St. Croix near Prescott, WI by launching boat at the public boat launch on the Mississippi River in Prescott, WI. Travel up the St. Croix River to sample site, conduct water quality sampling from boat. Collect an integrated sample using an integrating tube, collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel back to lab and process samples.
Significant Hazards as determined in column 3 of Hazard Analysis Form	(2) The St. Croix River in Washington County has been designated as infested with Eurasian watermilfoil by the MNDNR.	(2) Zebra mussels and rusty crayfish present
Limits for Each Control Measure	(3) No viable plant parts remain on boat, trailer, equipment, or waders.	(3) No zebra mussels or rusty crayfish remain on boat, trailer, equipment, or waders.
Monitoring Describe what is being monitored Explain how the monitoring will take place	(4) Presence of plant material. Ensure that boat, trailer, waders, and all equipment are free of viable plant parts. (5) Visual inspection of boat, trailer, waders, and other equipment for plant material.	(4) Presence of zebra mussels and rusty crayfish. Ensure the boat, trailer, waders, and other equipment are free of rusty crayfish and zebra mussels. (5) Visual inspection of boat, trailer, waders, and other equipment for rusty crayfish and zebra mussels.
Frequency of monitoring	(6) Each time, before and after boat, trailer, and equipment is used at site.	(6) Each time, before and after boat, trailer, and equipment is used at site.
Person or position responsible for monitoring	(7) Water quality monitoring staff.	(7) Water quality monitoring staff.
Corrective Actions Actions taken when limits of control measures are not met	(8) Cease operation and clean boat, trailer, waders, and equipment before launching onto water.	(8) Cease operation and clean boat, trailer, waders, and equipment before launching onto water.
Verification Method of verification	(9) Review records.	(9) Review records.
Records List what is recorded at each critical control point	(10) Record the procedure used to remove plant fragments. Record that inspections occurred prior to leaving one site and entering the next site.	(10) Record the procedure used to remove zebra mussels and rusty crayfish. Record that inspections occurred prior to leaving one launch and before entering the river at the next site.

This form accommodates two Critical Control points. Attach additional pages of this form as necessary.

Final Step: Once the HACCP plan is completed, attach it to the signed procedure form with the hazard analysis worksheets.

Once the HACCP plan is completed, attach it to the signed procedure form with the hazard analysis worksheets.

AIS-HACCP PLAN

Aquatic Invasive Species-Hazard Analysis and Critical Control Point AIS-HACCP Plan Form (continued)

Critical Control Point Each row answered "yes" in column 6 of Hazard Analysis Form	<i>(10) Day 3 continued: Conduct water quality monitoring on the Lake St. Croix near Prescott, WI by launching boat at the public boat launch on the Mississippi River in Prescott, WI. Travel up the St. Croix River to sample site, conduct water quality sampling from boat. Collect an integrated sample using an integrating tube, collect water quality data with a multiprobe and through field observations. Return to boat launch, trailer boat, travel back to lab and process samples.</i>
Significant Hazards as determined in column 3 of Hazard Analysis Form	<i>(2) The St. Croix River in Washington County has been designated as infested with Eurasian watermilfoil by the MNDNR.</i>
Limits for Each Control Measure	<i>(3) No viable plant parts remain on boat, trailer, equipment, or waders.</i>
Monitoring Describe what is being monitored	<i>(4) Presence of plant material. Ensure that boat, trailer, waders, and all equipment are free of viable plant parts.</i>
Explain how the monitoring will take place	<i>(5) Visual inspection of boat, trailer, waders, and other equipment for plant material</i>
Frequency of monitoring	<i>(6) Each time, before and after boat, trailer, and equipment is used at site.</i>
Person or position responsible for monitoring	<i>(7) Water quality monitoring staff.</i>
Corrective Actions Actions taken when limits of control measures are not met	<i>(8) Cease operation and clean boat, trailer, waders, and equipment before launching onto water.</i>
Verification Method of verification	<i>(9) Review records.</i>
Records List what is recorded at each critical control point	<i>(10) Record the procedure used to remove plant fragments. Record that inspections occurred prior to leaving one site and entering the next site.</i>

This form accommodates two Critical Control points. Attach additional pages of this form as necessary.

Final Step: Once the HACCP plan is completed, attach it to the signed procedure form with the hazard analysis worksheets.

Standard Operating Procedure #6: Field Measurements and Water Sample Collection

Version 1.2

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Revision History Log

The following table lists all edits and amendments to this document since the original publication date. Information entered in the log must be complete and concise. Users of this standard operating procedure will promptly notify the project manager and/or the Great Lakes Network (GLKN) data manager about recommended and required changes. The project manager must review and incorporate all changes, complete the revision history log, and change the date and version number on the title page and in the header of the document file. For complete instructions, please refer to Revising the Protocol, SOP #13.

Previous Version Number	Revision Date	Author (with title and affiliation)	Location in Document and Concise Description of Revision	Reason for Change	New Version Number
1.0	2/18/2010	Richard Damstra, David VanderMeulen, Joan Elias, GLKN	Clarified some verbiage throughout, made changes to field sequence, field gear checklist, added optical DO probe info, equipment blank procedures, and data backup procedures.	To clarify protocol and bring field procedures and protocol into agreement, based on new equipment and staff field experience.	1.1
1.1	2/01/2016	David VanderMeulen, Rick Damstra GLKN	Updated the NPS contact author and authorship, added information about continuous monitoring, fixed a handful of grammatical errors throughout	To clarify and update SOP based on changes that have occurred since Version 1.1 was published in 2010	1.2

*Add rows as needed for each change or set of changes tied to an updated version number

Acknowledgements

Combinations of existing protocols were used to develop this standard operating procedure for measuring the required water quality field parameters for the Great Lakes Network (GLKN). We consulted protocols written for the EPA, USGS, an online curriculum for monitoring water quality, this and other NPS networks, NPS internal documents, and the NPS–WRD website. We wish to acknowledge these sources and thank them for the valuable contribution that they have made to this document.

6.0 Introduction

Field measurements should represent, as closely as possible, the natural condition of the surface water at the time of sampling. Experience with and knowledge of the sampling equipment and the collection, storage, and processing of water samples for subsequent laboratory analyses are critical for collecting data of high quality. To ensure consistent, high-quality data, always:

- Make field measurements only with calibrated instruments that have been error-checked.
- Maintain a permanent log book for each field instrument for recording calibrations and repairs. Review the log book before leaving for the field. This book should also be used for recording results of pre-and post-calibration checks as well as housing copies of results from alternate measurement sensitivity (AMS) checks, if applicable.
- Test each instrument (meters and sensors) before leaving for the field. Become familiar with new instruments and new measurement techniques before collecting data.
- Have backup instruments readily available and in good working condition, whenever possible.
- Follow quality assurance/quality control procedures in SOP #12. Such protocols are mandatory for every data collection effort, and include practicing good field procedures and implementing quality-control checks. Make field measurements in a manner to minimize artifacts that can bias the result. Check field-measurement precision and accuracy (variability and bias).

6.0.1 Core Suite of Water Quality Variables

The core field variables are temperature, specific conductance, pH, water level, and dissolved oxygen, all of which are mandated by the National Park Service Water Resources Division (NPS-WRD). To this mandated suite, GLKN has added a measure of water clarity. Depth profiles of temperature, specific conductance, pH, and dissolved oxygen will be measured at each sampling station using a multiparameter instrument (multiprobe). Water clarity will be measured with a transparency tube. A Secchi disk will be used to collect additional water clarity measurements on impoundments or on other lake-like sites. Flow is a key parameter for physical conditions in rivers and is needed to interpret concentration data and calculate water quality loadings. Flow conditions will be tracked using the network of U.S. Geological Survey (USGS) gauging stations within Network large river parks. The expected ranges and resolutions required of multiprobes are listed in Table 6-1, and the methods for taking field measurements are described in Section 6.4.

6.0.2 Advanced Suite of Water Quality Variables

The advanced suite variables consist of total phosphorus, total nitrogen, nitrate+nitrite-nitrogen, ammonia-nitrogen, sulfate, chloride, magnesium, calcium, potassium, sodium, dissolved silica, dissolved organic carbon, alkalinity, total suspended solids (i.e., turbidity), and total chlorophyll-*a*. The methods for collecting water samples for these variables are described in Section 6.5. These water quality variables were chosen to monitor effects of the likely stressors on the resource, including land use change and atmospheric deposition. Other variables symptomatic of

contamination (e.g., DDT and PCBs) are addressed in GLKN's protocol for monitoring bioaccumulative contaminants (Route et al. 2009).

6.1 Equipment and Supplies

6.1.1 Instrument Specifications

Toward low bias and high precision of water quality data for this protocol, Table 6-1 lists typical field sensor performance specifications that should be expected from monitoring equipment for this protocol. Multiparameter sensor instruments (or multiprobes), while expensive, are convenient and commonly used in water quality monitoring. A multiprobe instrument should be selected based on the observable range for each variable. However, an additional investment in a back-up set of less expensive individual sensors will help ensure the development of a long-term data set, should the multiprobe not be working properly.

Table 6-1. Typical sensor performance specifications^a for field instruments (Penoyer 2003).

Sensor	Expected Range	Reporting Resolution ^b	Estimated Bias ^c
Temperature	-5°C to 45°C	0.01°C	±0.15°C
Specific Conductivity (SC25)	0 to 2000 µS/cm	1 µS/cm (range dependent)	±0.5% of reading + 1 µS/cm
pH	1 to 14 units	0.01 unit	±0.2 units
Dissolved Oxygen (Concentration)	0 to 50 mg/L	0.01 mg/L	0 to 20 mg/L ±0.2 mg/L; 20 to 50 mg/L ±0.6 mg/L
Dissolved Oxygen (% saturation)	0%–200%	0.1%	ca. ±2 %
Turbidity	---	---	---
Depth - Z (pressure sensor)	0→100 m	0.1 m	ca. 0.1 m

^a In the case of field probes, accuracy is typically a best-case maximum deviation from known correct values (typically based on comparisons with known NIST certified reference materials or standards).

True accuracy is a combination of high precision and low bias (see Irwin 2004 for more details).

^b Resolution/sensitivity is a data quality indicator related to detection limits but typically handled differently for field probes than for laboratory parameters. For more information, see Irwin (2004).

^c Estimated bias specifications reflect the uncertainty in measurement of the instrument and sensor in combination only, and not other factors that can affect accuracy, such as environmental variables or the ability of field personnel to calibrate and operate the instrument using proper protocols.

Instrument-specific estimations of the range of uncertainty for each variable must be calculated for use in the interpretation of data. Therefore documentation will be maintained regarding the ability of each multiprobe to meet the data quality objectives of this project; blank forms are included in SOP #12. The completed forms for each multiprobe will be maintained along with the calibration and maintenance logs for the multiprobe.

Details on estimating instrument sensitivity are included in SOP #12, Quality Assurance/Quality Control. Because field equipment is likely to change during the course of this long-term monitoring project, QA/QC procedures must be followed on documenting cumulative bias (SOP #12).

6.1.2 Additional Equipment, Supplies, Forms, etc.

Refer to the checklist of supplies and equipment needed for field sampling (Table 6-2) prior to each sampling trip. Keep on hand all necessary forms, calibration logbooks, field logbooks, field data sheets, procedural manuals, and equipment instructional manuals.

Table 6-2. Checklist of field equipment and supplies required for monitoring water quality.

√	Equipment and Supplies
	Field notebook, pencils, and pen (waterproof ink)
	New field forms on waterproof paper
	Up-to-date field folders containing recent data sheets for field comparison (copies only; never take originals in the field)
	Multiparameter instrument (calibrated), calibration standards, check solutions, data logger
	Long and short cables for multiprobe instrument(s)
	Calibration logbook for each instrument
	All maintenance parts and calibration standards for field instruments
	Backup instruments in case of electronic failure of multiprobe (for example: YSI 85 [T-DO-SC25] or equivalent, YSI 200 [T-DO], Hannah Dist3 [SC25], armored NIST certified thermometer [°C], portable pH meter)
	Transparency tube
	Secchi disk and metered line (marked at 0.5 m intervals)
	Sounding weight(s)
	Water samplers: Integrating tube, Van Dorn type sampler
	Compositing jug and other bottles, including brown or amber bottles for chlorophyll-a
	Sonde with barometric pressure sensor, weather radio barometer, or other way to obtain barometric pressure
	Pocket calculator (waterproof, if possible)
	Extra batteries for all field equipment (multiparameter probe, calculator, GPS, etc.)
	Rain gear
	First aid kit
	Personal flotation device(s)
	Field trip itinerary
	Cellular phone and/or park radio
	Digital camera with extra flash cards and battery
	Map(s) of station location, preferably at different scales
	Global positioning system (GPS) with extra batteries
	Deionized or distilled water for field rinsing

Copy field data sheets on waterproof paper. See the attachments for blank data sheets. The instruction manuals for each instrument should be copied and the originals placed in a secure file and kept in the office. Specific sections of the manual that might be important to have in the field should be copied onto waterproof paper and remain in the field kit. Include copies of a datalogger software manual.

6.2 Overview of Field Work

6.2.1 Sequence of Activities During Field Work Day

1. Review field gear checklist.
2. Create a new field form for each monitoring station, printed on waterproof paper.
3. Prepare sample bottles and labels in advance and place in a cooler.
4. Conduct daily calibration of appropriate meters and probes.
5. Inspect vehicles at the beginning of every field day, including all safety and directional lights, oil, gasoline, tire air pressure levels.
6. Inspect boat; ensure all safety gear is on board.
7. Drive to boat landing or sample site. If needed, load boat with sampling gear, launch boat, and navigate to monitoring site, otherwise walk to sample site. Set up a clean work space on the boat, or streambank for sampling.
8. Refer to description of monitoring station location, directions, maps, and photo to verify correct location. Verify coordinates on GPS unit.
9. Measure field water quality variables per Section 6.4 and collect samples per Section 6.5.
10. Be sure that all samples are correctly labeled and preserved on ice.
11. Verify that field form is completely filled out, and initial the form.
12. If sampling from more than one monitoring station in a day, follow procedures for decontamination of equipment per SOP #5, and go back to step 6, above.
13. Upon return to shore, inspect boat, trailer, waders, and all equipment that has come into contact with the water for invasive species.
14. Return to office or field station.
15. Process samples according to SOP #7. Refrigerate or freeze samples, as required and package samples for sending to contract analytical laboratory.
16. Clean sampling equipment per SOP #7. Rinse sensors with deionized water and perform calibration re-checks, as detailed in SOP #12. This can be done upon returning to home base. Perform decontamination of the probe or other instruments as detailed in SOP #5.
17. As soon as possible after returning from the field, review both hardcopy and electronic data; offload electronic data onto computer; review laboratory data as it is received.
18. At the end of the field season, enter or import all field and laboratory data into NPSTORET.

6.2.2 Recording Field Information Upon Arrival at Monitoring Site

Consistent methods are important to long-term data quality. In actuality, the ideal conditions are not always met in the field or in the lab and changes in staff occur. Therefore, documentation of procedures, site conditions, laboratory analysis, and reasons for deviations of any kind is important. Personnel are encouraged to write down more than they feel may be necessary in the moment, as the future interpretation of their data will depend on the written record and not the memory of an individual. Waterproof field forms (copy available in the attachments) should be prepared ahead of time labeled with the project and station IDs. Sampling stations will be identified by park and water body name according to GLKN guidance. Information on the sampling station and park will comply with NPSTORET and Aquarius database requirements. Field sampling forms are used to record the physical and chemical water quality variables measured at the time of sample collection. In addition to recording the field variables, any samples collected for laboratory analyses must be so indicated. Documentation should include calibration data for each instrument, field conditions at the time of sample collection, visual observations, and other information that might prove useful in interpreting these data in the future.

While at each monitoring site, the information recorded on field sampling forms should include:

- Date
- Time of arrival
- Names of field team members, with the person acting as crew leader listed first
- GPS coordinates, to verify location
- Current weather (air temperature and wind speed) and relevant notes about recent weather (storms or drought), including days since last significant precipitation
- Observations of water quality conditions (see below)
- Multiparameter meter (model/serial no.), calibration date, and field measurements of core suite variables; add dissolved oxygen calibration information if calibrating on-site
- List of sample IDs and collection times for advanced suite variables or quality assurance samples
- Whether any samples were not collected, and reason
- Quality assurance/quality control procedures followed
- Water level measurement, if collected at a given site
- Qualitative assessment of flow conditions (e.g., low, average, high)
- Any other required metadata for NPSTORET data entry
- Any additional non-routine monitoring (e.g., collection of dragonfly larvae for mercury analysis) by GLKN or park staff, or park partners
- Time of departure

All entries should be made clearly. If an incorrect entry is made, a single heavy line should be drawn through the incorrect entry and the correction made. All corrections should be initialed and dated. The completed field forms will be maintained in chronological order by station, copied into project binders and the originals maintained on file indefinitely. Field data are reviewed annually by GLKN personnel (see SOP #8, Data Entry and Management, for details).

Upon arrival at the monitoring station, record visual observations of water quality conditions that will be useful in interpreting water quality data.

- Water appearance—General observations on water may include color, unusual amount of suspended matter, debris, or foam.
- Biological activity—Excessive macrophyte, phytoplankton, or periphyton growth. The observation of water color and excessive algal growth is important in explaining high chlorophyll-*a* values. Other observations to note include types of fish, birds, or spawning fish.
- Unusual odors—Examples include hydrogen sulfide, mustiness, sewage, petroleum, chemicals, or chlorine.
- Watershed activities—Activities or events that are impacting water quality; for example, forest fires, road construction, timber harvest, shoreline mowing, beaver activity, or livestock watering.

6.2.3 Preventing Contamination

Field technicians should be aware of and record potential sources of contamination at each field site. Decontaminate field sampling equipment according to SOP #5 for minimizing the risk of spreading invasive species. Clean field and laboratory equipment according to SOP #7 to avoid contamination of analytes to be measured. Do not allow sample water to touch hands; do not touch insides of sampling equipment, containers, or laboratory bottles.

6.2.4 Using Disposable Gloves

Wearing disposable gloves is strongly recommended when handling acid preservatives. Check manufacturer's chemical resistance information to be sure gloves are appropriate for compounds to which they will be exposed. Common glove types include those made of vinyl, latex, and nitrile; nitrile is in standard use for USGS sampling work because of its resistance to most of the chemicals to which it typically will be exposed for the length of exposure (usually <15 minutes). Field personnel are cautioned that direct contact with materials such as latex or nitrile can cause severe allergic reactions in some individuals and should be monitored.

Physical properties to consider when selecting disposable gloves are glove length, slip protection, puncture resistance, heat and flame resistance, cold protection and comfort. These factors can vary between manufacturers. Gloves should be inspected visually for defects. Check for tears, punctures and other flaws that can prevent the glove from being an effective shield. After putting the gloves on, rinse them with water while gently rubbing hands together to remove any surface residue before handling sampling equipment.

6.2.5 Bottle Preparation – Types and Sizes of Bottles

For each monitoring station, select the bottles appropriate for each analyte and label them with Station ID, sample date, and analyte code according to the requirements of the contact analytical laboratory. Store pre-labeled bottles in a dry box or in separate bags for each station.

6.3 Field Measurement Procedures

6.3.1 Where and When to Measure Field Variables

The thalweg (deepest point in the riverbed) is the preferred sampling site for sites with flowing waters; for impoundments or other lake-like sections, the deepest area is sampled. If the morphometry is not known, sample at mid-stream unless there is another basis for selecting the site, such as high water making wading into the thalweg or mid-stream too dangerous. Upon visiting a site that has been visited before, a GPS should be used to locate the sample site. If a GPS is not available, use a compass and known landmarks to triangulate the sample site. A depth finder can also be used to verify that the deepest part of the river/impoundment is being sampled.

Discrete routine monitoring will typically occur eight times throughout the open-water season, April through November. This sampling timeline will provide ample sampling dates to be able to fully characterize the major period of plant/algae growth, oxygen depletion, water clarity, temperature change, maximum public use, and is likely to capture spring and/or fall runoff events in many years.

Sampling should be conducted during mid-day, from 10:00 AM to 3:00 PM, when possible. Water samples and Secchi depth readings are typically collected during mid-day to reduce variability in the data due to the differences in daylight, and to a lesser extent, temperature (Carlson and Simpson 1996). Mid-day is usually the peak period of algal photosynthesis, which can have dramatic effects on DO, transparency, and dissolved inorganic phosphorus (phosphate) and dissolved inorganic nitrogen (nitrate+nitrite+ammonium) throughout a 24 hr cycle.

For all sampling, it is critical to avoid sampling water showing evidence of oil, gasoline or anything else from the boat motor. It is best to turn off the engine and set the anchor, although this may not be possible or advisable in bad weather or with a balky engine. The engine is commonly located in the stern of the boat; therefore, the anchor should be secured near the prow of the boat, such that current or winds will rotate the stern away from the sampling area. After setting anchor, allow surface water to clear of disturbances. Collect samples on the upwind and/or upstream side of the boat, to minimize contamination and disturbance. Avoid surfactants, floating debris, and turbid aeration during sample collection. Discard rinse water or excess sampling water on the downwind/downstream side of the boat.

6.3.2 Stabilization of Sensor Probe Readings

Before making field measurements, properly-calibrated sensors must be allowed to equilibrate to the condition of the water being monitored. Sensors have equilibrated adequately when instrument readings have stabilized, that is, when the variability among measurements does not exceed an established criterion. The criteria for stabilized field readings were defined by O’Ney (2005) for a set of three or more sequential measurements (Table 6-3). Although the criteria used by the Greater

Yellowstone Network (O’Ney 2005) differ from those used in the upper Midwest by the University of Minnesota Natural Resources Research Institute (WOW 2004), the differences are small in comparison to the true range of variability one might expect in the field and with aging instrument sensors. The natural variability inherent in surface water or ground water at the time of sampling generally falls within these stability criteria and reflects the accuracy that should be attainable with a calibrated instrument. In the case of field probes, accuracy is typically a best case maximum deviation from known correct values (typically based on comparisons with known NIST certified reference materials or standards). True accuracy is a combination of high precision and low bias (see Irwin [2004, 2006] for more details).

Dissolved oxygen typically requires the greatest amount of time to stabilize. For dissolved oxygen sensors that utilize a membrane, differences in polarigraphic sensor membrane thicknesses, age, and rates of oxygen consumption increase the variability of the equilibration time. The longest equilibration times will typically occur where dissolved oxygen exhibits a steep gradient (change in DO concentration >5 mg/L) or very low oxidic levels (DO concentration <3 mg/L). Depending on the site characteristics and the specific oxygen sensor, 3 to 5 minutes may be required for complete equilibration. This time far exceeds what is needed for the other variables, which typically stabilize in less than 60 seconds. Observers should only note instrument DO readings after the stabilization criteria in Table 6-3 are met, and then record readings for all variables at once.

Table 6-3. Recommended instrument stabilization criteria for recording field measurements.

Standard Direct Field Measurement	Stabilization Criteria (O’Ney 2005)	Stabilization Criteria <i>In situ</i> Multisensors (WOW 2004)
Temperature:		
Thermistor Thermometer	± 0.2°C	± 0.2°C
Liquid-in-glass Thermometer	± 0.5°C	(5%)
Specific Conductivity (SC25)		
When ≤100 µS/cm	± 5%	< 5 µS/cm
When >100 µS/cm	± 3%	(10%)
pH: Meter displays to 0.01	± 0.1 unit	± 0.2 unit (10%)
Dissolved oxygen:		
Amperometric (same as polarigraphic) method	± 0.3 mg/L ± 2%	± 0.5 mg/L (10%)

6.3.3 Outline of Discrete Water Profile Measurements

Acquiring high quality results requires the use of consistent measurement methods. Adhere to the following guidelines:

1. Depths <20 m: Measure T, DO, pH, and SC25 surface to bottom at 1 m intervals.
2. Depths >20 m: Decrease measurements to 2 m intervals down to 30 m and to as much as 5 m increments for greater depths.

3. Wait for the DO value to stabilize first, record the value, then read the other parameters. Because DO takes the longest to stabilize this assures all parameters have equilibrated. See section 6.4.4 for further details. Stabilization of the DO value will typically take anywhere from 30 seconds to several minutes, depending on the gradient from the previous depth and the age and type of oxygen sensor. Extra time should be allowed for equilibration when values are below approximately 3 mg/L. If the sonde has a Clark cell DO sensor and does not have a stirring mechanism, jiggle the cable gently approximately once per second. Jiggling or stirring should not be needed for optical DO sensors.
4. Enter all data on field forms.
5. Quality Assurance (QA): At a minimum, replicate one of every 10 sets of measurements (e.g., at 1 m, 10 m, etc.). The replicate should be taken immediately following the original reading. Values should agree within + 10% or the detection limits listed in Table 6-3, whichever is larger.
6. Additional QA: At the first site sampled (once per field day at least), when a multiprobe is used to record water quality parameters, record 7 sets of measurements at the same depth after all parameters have stabilized. The readings should be taken immediately after one another in quick succession. These readings will be used to calculate the alternative measurement sensitivity (AMS) of each sensor on the multiprobe, which provides a way to track long-term sensor accuracy.

Instrument problems or failure

- If water quality sensor measurements are not representative of field conditions based on previous data or limnological knowledge, re-calibrate and try again.
- If readings seem reasonable, proceed. If not, first check the troubleshooting guide in the instrument manual. If problem persists, collect as much data as possible using back-up hand held instruments, if available. Using the Van Dorn bottle, collect a sample from 2.5, 5, 7.5, and 10 m, and then at 5 m intervals to the bottom, and record values obtained by dipping the sensors into the top of the sampler. In the case of measuring DO, jiggle the probe without causing bubbles to form in low DO hypolimnetic samples. If, after approximately 2 min, the value continues to decrease, increase the rate of swirling or jiggling to see if it will stabilize; if it is increasing after 2 minutes, you may be agitating the water enough to be causing an aeration artifact near the surface. If too much time elapses and the temperature of the water sample has increased by more than 1°C, collect a fresh sample.

6.3.4 Detailed Description and Troubleshooting Hints for Field Variables

Because temperature, DO, and other water quality variables are important determinants of biotic habitats, it is important that observers write down values on field forms and think about their ecological meaning, even if a data logger is recording the measurements. The hard copy also serves as backup in case there is an electronic failure.

6.3.4.1 Temperature

Temperature (T) is measured in units of degrees Celsius (°C) and recorded to the nearest degree or tenth of a degree as warranted by instrument.

1. If a cabled thermistor is not available, the high specific heat of water allows a temperature profile to be obtained by bringing deep samples to the surface with a water sampler and immediately measuring temperature with a hand-held thermometer or thermistor. Anything immersed into the water sampler potentially contaminates that particular sample; a separate sample must be collected for water chemistry.
2. The upper few centimeters of soft sediments are often several tenths of a degree warmer than the overlying water. The rise in temperature can be an indication that the probe is submersed into the sediments. If this happens, be sure to vigorously shake the instrument in shallow water with high DO to clean it before re-taking measurements. Check intermediate depth values, and if these values do not meet QA criteria, pull the instrument to the surface and clean it.

6.3.4.2 Specific Conductivity

Specific conductivity (SC) is the ability of water to conduct an electrical current for a unit length and unit cross-section at a certain temperature, measured in units of microsiemens per centimeter ($\mu\text{S}/\text{cm}$), and recorded to the nearest $\mu\text{S}/\text{cm}$. Commonly used in water quality monitoring, SC is a general measure of the number of ions dissolved in the water. It is important to be aware of the difference between SC (specific conductivity at the ambient temperature of the sensor) and SC25 (an abbreviation for specific conductivity temperature compensated to 25°C). This difference becomes very important in profiling stratified areas where the hypolimnion is cold with respect to the epilimnion. In such cases, the typical increase in ions with depth, as estimated by SC25, would be substantially offset by the corresponding decrease in SC due to decreasing temperature. The difference between SC and SC25 can confound analyses of seasonal patterns of dissolved ions since water temperatures vary throughout the year. The SC25 can be used to monitor seasonal changes in total dissolved salts (TDS) such as a spring flush of road salt, which is why the temperature compensation is so important. Many instruments will display SC in addition to SC25. In the event that an uncompensated sensor must be used (older instruments and most pocket conductivity meters are not temperature-compensated), the value of SC25 can be calculated from SC and temperature values (see # 3 below). The value of SC should be recorded on the field form even when SC25 will be calculated.

1. A common physical problem in using a specific conductance probe (or meter) is entrapment of air in the conductivity probe chambers. Its presence is indicated by unstable specific conductance values fluctuating up to $\pm 100 \mu\text{S}/\text{cm}$. This problem is much more prevalent in turbulent stream waters and can be minimized by slowly and carefully placing the probe vertically into the water and when completely submerged, quickly moving it back and forth through the water to release any air bubbles. An SC probe with an open flow design does not trap air.

2. Is the value real or is the instrument out of calibration? Having specific conductance standards in the field can help verify values that fall outside the expected range. For example, the expected specific conductivity is around 200 and the reading is 1500. A known standard can be put in the instrument storage cup to determine if the instrument is reading correctly or is out of calibration.
3. SC25 values can be calculated from uncompensated SC values via a temperature-compensation formula. For example, for YSI probes, use:

$$SC_{25} = \frac{SC_m}{1 + 0.0191(t_m - 25)}$$

Where, SC_{25} = corrected conductivity value adjusted to 25°C,
 SC_m = measured conductivity before correction, and
 t_m = water temperature at time of SC_m measurement.

Contact the instrument vendor for the appropriate formula.

6.3.4.3 Hydrogen Ion Activity (pH)

Commonly used in water quality monitoring, pH is a measure of the acidity of water, measured in standard pH units (SU), and recorded to the nearest 0.1 pH unit. The pH scale is from 1 to 14: neutral water is pH 7, acidic waters have pH <7, and alkaline waters have pH >7.

- Is the value real or is the instrument out of calibration? Having pH standards in the field can help verify values that fall outside the expected range. For example, the expected pH is around 7.0 and the reading is 9.5. A known standard can be put in the instrument storage cup to determine if the instrument is reading correctly or out of calibration.
- As with dissolved oxygen, a pH probe can take longer to equilibrate when the gradient from the previous measurement is large (>1.0 pH SU).
- Low ionic strength waters with SC25 <50 µS/cm can cause pH measurement stability problems with some probes, necessitating use of low ionic strength probes. Probes will often calibrate fine in strong ionic strength buffers but will not read accurately in lower ionic strength surface waters. If you suspect this is the case, use a sensor that is designed for low ionic strength waters.
- Because the pH scale is logarithmic, field values may first have to be converted to hydrogen ion activity (i.e., concentration) values, averaged, and re-transformed to pH standard units prior to conducting statistical analysis. To compute a mean pH for a group of data:
 - Convert each pH value to hydrogen-ion activity, using the equation: Hydrogen Activity = $10^{-\text{pH}}$.
 - Calculate the mean of the activity values by adding the values and dividing the sum by the total number of values.
 - Convert the calculated mean activity back to pH units, using the equation, $\text{pH} = -\log_{10}(\text{mean hydrogen activity})$.

6.3.4.4 Dissolved Oxygen

Dissolved oxygen (DO): units of mg/L record value to nearest 0.1 mg/L unless otherwise justified; percent saturation (% DO) record value to nearest %; also temperature compensated to 25°C.

- Be aware that if a water sample has a strong rotten egg smell (H_2S gas) it must have a DO of zero. This is one way to check your meter. You can use the measured offset value to correct your higher DO values. Do not report negative DO values in the final database although it is important to report them on the field data sheet. Do the correction afterwards but note on the field sheet the depth at which you could smell sulfide gas. Avoid touching the bottom, if possible, as the sensor may become fouled.
- Equilibration time is critical; the steeper the DO gradient, the longer the equilibration time. It may take >5 minutes when DO drops abruptly to near zero.
- The DO probes with membranes (Clark cell) actually consume oxygen in the immediate vicinity around the membrane as they work; measurements therefore require moving water using either a built-in stirrer (typical in multiparameter sondes and BOD probes) or moving the cable up and down (e.g., 6" each side of the desired depth) during the measurement. Optical sensors do not consume oxygen and hence do not require moving water.\
- Accuracy of an optical DO probe can be compromised if it is covered with a residue that inhibits or increases oxygen reaching the sensor surface. Algae on the sensor surface may increase DO measurements, while oils or sediments may lower them. If measurements seem suspect, or if the instrument was used in contaminated water, the sensing surface should be cleaned with a soft brush and a mild detergent.
- If the electronic DO meter is not functioning properly, DO can be measured by Winkler titration (Clesceri et al. 1998). A variety of field kits are convenient and cost effective, including Lamotte, Hach, and others.

6.3.4.5 Clarity

Measurement of Secchi disk transparency has historically been the most common means of measuring water clarity due to its simplicity. It is a qualitative evaluation of the transparency of water to light based on the reflection of light from the surface of the Secchi disk and is a function of the absorption characteristics both of the water and its dissolved and particulate matter. To get an accurate measure of Secchi depth, the disk must hang vertically from the side of the boat. The transparency tube, which uses the same principle as a Secchi disk, is recommended when the Secchi disk can be seen on the bottom or when the disk does not hang vertically due to a current or an unstable boat.

6.3.4.5.1 Secchi Depth:

1. Measure Secchi depth using a 20 cm (8 in diameter) black and white Secchi disk from the shaded side of the boat. Do not wear sunglasses while viewing the Secchi disk.
2. Measurements should be made as near to midday as possible (10 AM to 3 PM; sunny and calm is optimal). Visit each site as close to the same time of day as previous visits in order to minimize diurnal variation.

3. Try to lower the disk in the shade of the boat and make sure that the line is hanging vertically. In strong currents or choppy waves, it may be helpful to tape a rock or weight to the bottom of the disk. If sampling an area with a discernable current, a Secchi measurement is not needed.
4. Slowly lower the disk into the water until it disappears and note the depth.
5. Lower the disk a little farther, then slowly raise it until the disk reappears and note the depth again.
6. Average the two readings and record the value. Record the Secchi observer on the data sheet (see #8 below).
7. Record the conditions present when the Secchi measurement was made, with 1 being excellent (sunny clear skies, calm water), 2 being moderate (some clouds, small waves), or 3 being poor (dark skies, very choppy water, or currents that make it extremely difficult to read).
8. Quality Assurance (QA): At a minimum, replicate Secchi readings at every tenth site. Because of the apparent ease of this measurement yet its potential difficulty in less than ideal conditions, all field crew members should take this measurement at each site for at least the first round of sampling. The crew should not reveal their value until all are finished and then all values should be recorded and compared. Values should agree within $\pm 10\%$ for Secchi measurements < 5 m and ± 0.5 m for greater values.

6.3.4.5.2 Transparency Tube:

1. Measure transparency tube clarity using a 120 cm tube at all sites. For all shallow (< 2 m) sites or sites with discernable current, measure clarity with a transparency tube even if Secchi measurements can be made on occasion because clear water times can occur, when Secchi depth cannot be measured.
2. The tube should be set on a white towel background, shaded by your body, and read without sunglasses.
3. Water should be dispensed from a carboy that was used for integrating water samples (see below) rather than dipped from the water body. The carboy must be well shaken prior to filling the tube to minimize artifacts due to settling of sand and larger silt particles. Allow air bubbles to disappear before making the final measurement.
4. While slowly releasing water from the bottom of the tube via its valve, note and record the depth at which the mini-Secchi first becomes visible. Discard the remaining water and repeat the measurement with a second subsample from the carboy.
5. Several attempts to read the clarity may be necessary because of overshooting the endpoint, so collect plenty of water for this analysis. A standard 120 cm $\times$ 4.5 cm outside diameter tube requires approximately 1.5 L to fill it, so dedicate at least 4 L of water for this measurement.
6. During clear water periods, the tube may not be long enough for a measurement. In such cases, the value should be recorded as > 120 cm. If it appears to be barely visible, this fact should be recorded to distinguish it from a measurement where it is clearly visible.

7. Quality Assurance (QA): At a minimum, replicate transparency tube readings at every tenth site. As for the Secchi reading, because of the apparent ease of this measurement yet its potential difficulty in less than ideal conditions, all field crew members should take this measurement at each site for at least the first round of sampling. Crew members should not reveal their values until all are finished and then all values should be recorded and compared. Values should agree within 10%. This acceptance criterion is subject to change as more data from various volunteer monitoring programs become available (such as the Minnesota Citizen Stream Monitoring Program (<http://www.pca.state.mn.us/water/csmp.html>)).

Maintenance notes and other precautions:

1. The rubber stopper (with attached Secchi) can be dislodged easily. Tape the stopper with black vinyl electricians tape and carry an extra stopper-Secchi.
2. Clean the transparency tube periodically with mild dish soap and a soft cloth.
3. Although water from the tube may be saved for turbidity and TSS measurements, do not save it for nutrient or other pollutant analyses because the tubes are not cleaned according to certified protocols.
4. Subsampling and settling issues are important, as particles settle quickly. A stopper for the top of the tube is useful to allow for resuspension during the measurement if rapid settling occurs.
5. Dissolved color due to organic matter (humic and fulvic acids, usually from bogs and conifer needles) can confound comparisons of transparency tube data between sites. Also, areas with similar concentrations of suspended sediments can have different transparency because smaller particles scatter more light.
6. A transparency reading taken from one tube cannot be compared with a reading taken from another tube made by a different manufacturer if the dimensions are different.

6.4 Water Sample Collection

In flowing sites, water samples will usually be taken using an elbow deep grab sample, since in most cases these flowing waters will be well-mixed. When sampling off bridge deck sites, use a Van Dorn or Niskin type remote-closing water bottle to collect a grab sample from just below the surface of the water, and retrieve back up to the bridge, and fill the composite jug from the remote-closing bottle. Precautions will be taken to check with the manufacturer regarding the potential for nutrient or other contamination from the sampler. Before its first use, the sampler will be cleaned thoroughly by rinsing three times with hot tap water, rinsing three times with 0.1 N HCl, and finally, rinsing three times with deionized water.

In lake-like sites, a tube sampler collects and integrates (i.e., an integrated sampling tube or I-tube) a column of water into a single sample. If the top end is capped, all or most of the water will remain in the tube as it is raised. The main advantage over using a surface dip (often called a grab sample), is that a larger stratum of water in the epilimnion is represented by the sample, reducing the effect of

high chlorophyll concentrations from concentrated scums on the seasonal mean. Bottom water, and on occasion other depths, will be sampled using a Van Dorn or Niskin type remote-closing water bottle. Although less durable, a transparent acrylic bottle allows the crew to ensure that near bottom water samples do not contain suspended bottom sediment before pouring the (contaminated) water into the sample bottles. Precautions will be taken to check with the manufacturer regarding the potential for nutrient or other contamination from the sampler. Before its first use, the sampler will be cleaned thoroughly by rinsing three times with hot tap water, rinsing three times with 0.1 N HCl, and finally, rinsing three times with deionized water.

6.4.1 Integrated Sampling Tube and Elbow-Deep Grab Samples

1. Rinse compositing jug 3 times with native water before sampling. Increase surface water flushing to 6 rinses if the compositing jug previously contained water that was contaminated with sediment or water deemed to have much higher nutrient levels.
2. If using I-tube, remove both stoppers from integrated sampler. Rinse the integrated sampler in native water 3 times (lower to 2 m, raise, lower again, etc. for 3 rinses).
3. If collecting I-tube sample by lowering sampler into water column (if water depth will allow it). If the site is too shallow for the 2 m sampler, then use one of the following methods:
 - a. collect sample by using the integrated sampler on an angle instead of vertically,
 - b. fill jug with 0.5 m deep water using the Van Dorn sampler,
 - c. or fill jug via surface grab.

Secure the top stopper, raise the sampler vertically, and immediately drain it into the rinsed compositing jug. The sampler will start releasing its water as soon as it clears the water surface. Two or three integrated samplers of water will be needed to fill a 6 L jug. Fill the jug at least 75% full to ensure adequate water for all analyses; if a transparency tube measurement will be done, an entire jug is needed. Extra water simply reduces variance associated with the site.

4. If collecting elbow-deep grab sample, lower the capped compositing jug under the water. Carefully uncapped while under water and allow the jug to fill completely. Replace the cap while still under water, and bring to the surface. Dispense water needed for chlorophyll-a analysis into appropriate bottle, and refill composite jug as needed using above procedure.
5. Once water is collected, immediately begin dispensing it into appropriate analyte bottles or keep jug cold and dark until processing in the park lab or home base at the end of the day (details below).

6.4.2 Discrete Depth Sampling

During the mid-summer sampling when lake-like areas are thermally stratified, (defined as a gradient of $\geq 1^{\circ}\text{C}/\text{m}$ or DO levels $\leq 2 \text{ mgO}_2/\text{L}$), collect two separate samples.

Collect surface water with the 0–2 m integrated tube sampler as above; collect bottom water with Van Dorn sampler from approximately 0.5 to 1 m above the bottom. Because the hypolimnetic water will be used for nutrient analyses only, a smaller container (1 or 2 L) may be used for storage and

transport to the sample processing site. Do not collect bottom water from sites that are not thermally stratified, as evidenced by temperature (and oxygen) profile data recorded with a multiprobe.

6.4.3 Distributing Sample Water from the Compositing Jugs

- Always ensure composite water is well mixed prior to dispensing it to any other containers.
- Rinse bottles and caps for chlorophyll-*a* with native water three times, prior to filling. Uncap and re-cap bottles below water surface to avoid surface scum or debris.
- Fill dark plastic bottle for chlorophyll-*a* (at least 1 L); keep cold and dark until filtering at the end of the day.

If the water sample is processed on-site, follow the instructions below, in addition to those in SOP #7.

Integrated 0–2 m or Elbow-Deep Grab Water

1. Rinse caps and bottles with composite water three times, using approximately 50 mL for each rinse, prior to filling. Shake out excess water.
2. Cap and re-agitate composite jug and then dispense water into previously labeled appropriate bottles. Be careful not to overfill bottles that already contain an acid preservative. Fill nutrient bottle to the bottom of the neck of the bottle – this will prevent the bottle from breaking when the water expands as it freezes.
3. Use remaining water for transparency tube measurement if specified. If insufficient water is left, collect additional water until there is enough for replicate measurements.

Bottom Water

- *From compositing jug*—Rinse the bottle caps and bottles in composite water as for surface water.
- *From Van Dorn*—Carefully stand the Van Dorn bottle on the side of the boat or on a seat, crack the top slightly to start the flow, rinse and dump 3 aliquots before filling the bottles to their necks. If there is not enough water in the Van Dorn to fill all the bottles, start over and either be more cautious with draining the sampler or use the compositing jug after rinsing, as above. Two Van Dorn samples may be pooled. Cap loosely and agitate the jug before dispensing water into the plastic bottles for raw and filtered nutrients (probably the only analyses to be done for bottom water).

Pay special attention to the appearance (visual color and turbidity) and smell (rotten egg gas, H₂S) of the water. If there is any evidence suggesting that bottom sediments were stirred up and captured by the sampler, re-do the collection taking care to vigorously clean the sampler and compositing carboy with surface water.

6.4.4 Sample Handling While in the Field

Store water samples in cooler with ice until return to home base for further processing.

In bad weather on sites with a single sampling station or with a small boat it will be more convenient and safer to sample the bottom water first, collect the surface sample after the profiling is done, and then return to shore to pour the subsamples from a full compositing jug.

6.4.5 Quality Assurance of Field Duplicates

Collect a field duplicate every 10 samples. Label the duplicate analysis bottles with the code appropriate for the duplicate, using the same date and time of sampling. Indicate on the data field form which site or is the duplicate. Treat the duplicate as a regular sample for all phases of collection, processing, and analysis. The duplicate should be a split sample, taken from the same composite jug. See section 6.6.3.2, below, for more details.

6.5 Departing From the Monitoring Site

Before leaving the monitoring site, all field forms and sample labels must be reviewed for legibility, accuracy, and completeness. Any changes in procedure due to field condition must be explained in the comments section. Make sure the information is complete on all forms. Record the departure time on the field form. After reviewing each form, initial the upper right corner of each page of the form. Document any photos taken by including the photo number and roll number or digital camera photo number on the field form.

6.6 Quality Assurance/Quality Control (QA/QC)

QA/QC basically refers to all those things good investigators do to make sure their measurements are right-on (accurate; the absolute true value), reproducible (precise; consistent), and include good estimates of uncertainty. It specifically involves following established rules in the field and lab to assure everyone that the sample is representative of the site; free from outside contamination by the sample collector, and that it has been analyzed following standard QA/QC methods. See SOP #12, Quality Assurance/Quality Control, for more details.

6.6.1 Calibration of Field Instrument Sensors

Calibration schedules overlap but differ from sampling schedules, so calibration methods are listed here as a separate procedural step. Instrument calibration is an essential part of quality assurance. Table 6-4 summarizes the ideal calibration frequency and minimum acceptance criteria for some of the most commonly collected water quality parameters. The reality of logistical constraints at back country sites may preclude calibration and checks of calibration at the ideal frequency. This SOP provides only generic guidelines for equipment use and maintenance. A wide variety of field instruments is available; such instruments are continuously being updated or replaced using newer technology. Keep equipment manufacturers' maintenance and calibration instructions for all instruments for reference purposes. Field personnel must be familiar with the instructions provided by manufacturers. Contact manufacturers for answers to technical questions.

Table 6-4. Ideal calibration frequency and acceptance criteria.

Parameter	USEPA Method	Minimum Calibration Frequency and QC Checks	Acceptance Criteria	Corrective Actions
Temperature	170.1	Annually, 2-point check with NIST thermometer	± 1.0 °C	Re-test with a different thermometer; repeat measurement.
Specific conductance (SC25)	120.1	Daily, prior to field mobilization; calibration check prior to each round of sampling; 10% of the readings taken each day must be duplicated or a minimum of 1 reading if fewer than 10 samples are read.	$\pm 5\%$	Re-test; check low battery indicator; use a different meter; use different standards; repeat measurement
pH	150.1	Daily, prior to field mobilization (two buffers should be selected that bracket the anticipated pH of the water body to be sampled)	± 0.05 pH unit	Re-test; check low battery indicator; use different standards; repeat measurement; don't move cords or cause friction/static
		Calibration check w/ third buffer prior to each round of sampling; check with low ionic strength buffer in addition, if conductivity is < 50 $\mu\text{S}/\text{cm}$	± 0.1 pH unit	
		10% of the readings taken each day must be duplicated or a minimum of 1 reading if fewer than 10 samples are read.	RPD 10%	
Depth	--	Daily, prior to field mobilization, check at the field site. Check annually against commercially purchased brass sash chain labeled every 0.5 m to ensure that it reads zero at the surface and varies < 0.3 m for depths < 10 m and no more than 2% for greater depths.	± 0.1 m	Retest, check low battery indicator; repeat measurement; use with accurately calibrated line

Table 6-4 (continued). Ideal calibration frequency and acceptance criteria.

Parameter	USEPA Method	Minimum Calibration Frequency and QC Checks	Acceptance Criteria	Corrective Actions
Transparency tube	--	Transparency tubes have a 100 or 120 cm scale; ensure tube is clean	±1.0 cm for transparency tube	
Marked lines (e.g., Secchi, Van Dorn)	--	Check markings annually against brass sash chain. If lines are heated (for decontamination) check prior to each round of sampling.	±1%, 0–10 m ±2%, >10 m	Re-mark line.

6.6.1.1 Instrument Calibration and Maintenance Logbooks

Calibration and maintenance logs for multi-parameter sondes and all back-up sensor probes will be maintained and will document the frequency of calibration, maintenance, and calibration checks. See the attachments for a blank calibration log form. Keep calibration logs with each instrument during the sampling season. Logs will later be archived at the Network office in Ashland, Wisconsin. A new log will be started for each field season. Each instrument will have a logbook for recording all maintenance and calibration information, including:

- serial number, date received, manufacturer's contact information, especially technical service representatives
- service records, dates of probe replacements
- maintenance records, for example, whenever the following general maintenance occurs: DO membrane replacement, pH reference probe junction and filling solution, probe cleanings, sonde (the sensor housing) replacement, impellor replacement or cleaning, etc.
- calibration dates and calibration data
- any problems with sensors
- pre-mobilization, post-calibration checks performed on individual sensor probes

6.6.1.2 Handling of Calibration Standard Solutions

Store all calibration standards in a temperature-controlled environment. Standards should be dated upon receipt and upon opening. Commercially-purchased calibration standards come with an expiration date that must be observed. Ensure that calibration standards are not used beyond expiration dates.

Properly dispose of all waste materials. Used calibration solutions, in general, may be rinsed down a sink with water after consideration of the wastewater treatment system available to that sink. Safety data sheets (SDS) that are sent with manufacturer purchased calibration solutions should be kept on file. These documents describe the flammability, toxicity, and other safety hazards of reagents. Some

reagents may include constituents that are toxic to aquatic life. These should not be rinsed down a sink in any large quantities in primitive areas where the ultimate destination of wastewater is the aquatic environment. Instead, these reagents should be collected in a properly-marked leak-proof container for disposal in an adequate treatment system.

6.6.1.3 Temperature

Temperature is typically not adjustable on an electronic sensor but should be cross-compared to a National Institute of Standards and Technology (NIST) traceable thermometer at the beginning of each field season, as follows:

- Compare against a NIST-certified or NIST-traceable thermometer at a broad range of temperatures, for example 0°C to 40°C;
- The sensor should read within $\pm 1.0^\circ\text{C}$ of the NIST thermometer;
- Typically you cannot adjust the instrument to calibrate it but check the manual. It is a good idea to check the instrument at 0°C in slurry of ice-water if a calibrated (NIST) thermometer is not available since electronic and non-electronic temperature sensors are typically linear over the likely range of field temperatures.

An armored glass thermometer that has been referenced to the NIST standard should be taken into the field for air temperature and surface water temperature measurements and for checks of the electronic sensor.

6.6.1.4 Specific Conductivity

Specific conductivity (SC25) will be calibrated using a KCl solution as specified by the instrument manual. Stock calibration solutions can be purchased commercially, prepared by a water quality contract lab, or made in an academic or agency lab. If the KCl solution is prepared by an academic or agency lab, it should be cross checked by an external certified lab. Set the instrument to record temperature-compensated SC (SC25) rather than SC.

Because the typical modern SC25 sensor is linear to <3% over the range from approximately 20 to 10,000 $\mu\text{S}/\text{cm}$, a single point calibration is typically sufficient. A typical standard is 1000 or 1413 $\mu\text{S}/\text{cm}$. Pre-mobilization error checks of this sensor using 10, 100, 1413, and 10,000 $\mu\text{S}/\text{cm}$ standards may be used to establish sensor error over the range of most interest in freshwater work.

6.6.1.5 pH

The pH is calibrated using the standard two buffer technique, using either pH 4 and pH 7, or pH 7 and pH 10, depending on the expected field values. Calibrate the probe according to the manufacturer's recommendations, usually starting with pH 7 buffer followed by the second buffer. If a water body is classified as low acid neutralizing capacity (ANC) acid-sensitive (i.e., ANC approximately 100 ueq/L or lower), it should also be checked against a low ionic strength buffer (LISB) with pH approximately 4, as per protocols from the National Acid Precipitation Assessment Program (NAPAP 1990). A low ionic strength pH combination electrode may be necessary to acquire this extra level of sensitivity if stabilized pH measurements are not achieved with standard

pH sensors. Prior to each round of sampling, check the calibration with a third standard with a pH value between those used for calibration.

6.6.1.6 Dissolved Oxygen

Two types of dissolved oxygen (DO) sensors are typically utilized: Clark cell and optical. A Clark cell (membrane) sensor is air-calibrated, while an optical DO sensor is calibrated in 100% saturated tap water. For a Clark cell, it is assumed that the dry sensor will read 100% saturation in an enclosed airspace with enough water in the bottom of the container to saturate the air with water vapor. For optical DO sensors, the sensor window must be submerged in 100% DO saturated water, which is obtained by agitating tap water in a container for one minute and then decanting it into the calibration chamber until the DO sensor is completely covered with water.

A Clark cell should be checked for bubbles under the surface of the membrane. If bubbles are observed, change the electrolyte solution and replace the membrane. An optical DO sensor's surface should be observed to make certain that no air bubbles are clinging to it. The calibration chamber can be tapped gently to dislodge any bubbles.

Temperature affects DO saturation, as does the air pressure, which varies with elevation and ambient weather. Both sensors require a known barometric pressure (BP), obtained by the multiprobe itself (if equipped with a BP sensor) or from another source. Acceptable sources for BP are from another instrument, such as a calibrated barometer, or a third-party source, such as the local weather bureau or airport. Typically, one can assume the barometric pressure to vary with elevation. As a result, the instrument should be re-calibrated at each site if the elevation has changed more than 50 feet.

6.6.1.7 Depth

The length measurement of brass Secchi chains, transparency tubes, and depth sounders usually require little calibration after purchase and an initial check for accuracy. Compare all depth cables to each other and ensure they are synchronized. A brass Secchi chain can be used to calibrate other cables and lines. The use of a brass chain is particularly important because this single tool is likely to provide consistent long-term data. Nylon and certain braided weave ropes and cords can stretch as much as 20% and may vary depending upon wetness, load, and age. Lines may also shrink, as when they are soaked in hot water to avoid spread of exotic species.

Electronic depth sensors, or pressure transducers, are based on pressure differences and need to be set to zero at the water surface initially. Perform this zero check at every station; however, verify the depth measurement with labels on the actual instrument cable.

A useful and widely-used labeling system for coding depths is:

1. 5 m increments: Wrap vinyl tape of a specified color at 5 m increments and add an extra wrap every 5 m. Therefore, there is a single wrap at 5 m: two wraps at 10 m separated by approx.. 0.5 cm with 10 m lying exactly between the wraps; three wraps at 15 m with the 15 m depth located at the middle of the second wrap, and so on. A permanent marking pen should be used to write the actual depth on the tape.

2. 0–10 m depth: Use a second color to make wraps at 1 m intervals to best define shallow water columns.
3. 10–20 m depth: Use the second color, or new third color, to wrap at 2 m intervals.
4. >20 m: Mark the line at 2.5 m intervals or continue with 2 m intervals.

6.6.1.8 Post-Field Calibration Checks

Post-field calibration checks must be performed after each use of the instrument and before any instrument maintenance. The sooner this procedure is performed, the more representative the results will be for assessing performance during the preceding field measurements. Calibration and post-calibration should be no more than 24 hours apart. When sampling daily, the second day's calibration can serve as the first day's post-field calibration check. Take the same care used in performing the initial calibration by rinsing the sensors and waiting for sensors to stabilize. After making measurements at the last station, fill the sampling cup with ambient water (not deionized or tap water). Repeat the initial calibration procedures performed before the sampling trip. Record post-field calibration values in the calibration logbook (generally on the same page with the initial calibration for that sampling trip). Deviation beyond the manufacturer's specifications is cause for concern and should be addressed before the next sampling date.

Do not adjust the instrument (using calibration controls) during the post-calibration check.

The purpose of the post-calibration is to determine if the instrument has held calibration during the day of sampling. Compare the post-calibration values to the expected values for the standards. This will ensure that the field measurements for the day can be reported with confidence. The difference between the post-calibration value and expected standard value can be used to indicate both calibration precision and instrument performance.

If post-calibration values (Table 6-5) fall outside the error limits for DO, pH, and specific conductance, data collected do not meet quality assurance (QA) standards and should be flagged appropriately (see SOP #12 for more details on QA/QC). Measurements may be repeated with a different or back-up instrument. If post-calibration measurements do not consistently fall within the error limits after in-house trouble shooting, the instrument should be returned to the manufacturer for maintenance.

Table 6-5. Post-calibration check error limits.

Parameter	Value
Temperature	±1°C, annual calibration check
Specific Conductance	±5%
pH	±0.1 standard units
Dissolved Oxygen	±0.2 mg/L, ± 10% saturation

6.6.1.9 Sensor Maintenance and Storage

Most multiparameter sondes should be stored with a small amount of water in the storage cup. Refer to the manufacturer's manual for tips on cleaning the probes and housing, routine maintenance procedures, and proper storage procedures.

6.6.2 Calculation of Field Instrument Performance Criteria

Performance of field instruments must be checked in several ways, as detailed in the QA/QC procedures of SOP #12. Formulae for calculating instrument sensitivity, instrument precision, instrument bias, and cumulative bias are found in SOP #12.

Documentation of these performance criteria include estimation of the limits of detection of the multiprobe, the relative percent difference of duplicates, measurement of reference solutions, and estimates of change due to change in methods or instrumentation.

6.6.3 Quality Assurance/Quality Control Samples

Quality control samples are commonly used in documenting quality control associated with the collection of samples in the field. Field blanks and duplicates are routinely incorporated into a monitoring program without a good understanding of their function. The purpose of these samples is to validate the precision and accuracy of laboratory data, and to determine the adequacy of preservation techniques, equipment cleaning and preparation, and sampling procedures. Field blanks are used to measure and quantify the amount of contamination from extraneous sources (preservatives, sample bottles, sample handling, automatic samplers, etc.) that might compromise the integrity of a sample (alter its true value or concentration).

6.6.3.1 Equipment Blanks

Blanks are an integral part of quality control (QC) and are required for all sampling activities. Their creation should be noted in the field log book. Blanks establish that there is no sample contamination from the containers during custody, transportation, and or pre-analysis preparation either in the field or in the laboratory. Blanks establish the level of constituents introduced into a sample by the equipment used for sampling, preservatives, and/or containers. We will conduct equipment blanks prior to each field season and occasionally during the field season between field sites to ensure field rinsing is adequate.

Collect an equipment blank prior to the field season as follows:

1. Clean all equipment used to collect, store, and process water samples (e.g., integrating sampler, compositing jug, filtering apparatus) according to SOP #7.
2. Rinse the equipment used to collect and store water samples (integrating sampler and compositing jug) with laboratory reagent grade water three times and discard.
3. Fill the integrating sampler with a fourth aliquot of laboratory reagent grade water and handle and process this aliquot as if it were a river sample to be analyzed.

Another source of systematic error is sample cross contamination from field sampling equipment used to handle a multiple number of samples. The integrated sampler and compositing jugs are to be

rinsed in the field three times at each site prior to taking a sample. Either piece of equipment may be a source of cross contamination. Collect periodic equipment blanks as follows:

1. In between sample sites, rinse the equipment used to transfer water samples (integrating sampler and compositing jug) with laboratory reagent grade water three times and discard.
2. Fill the integrating sampler with a fourth aliquot of laboratory reagent grade water and handle and process this aliquot as if it were a river sample to be analyzed.

This sample is labeled as an equipment blank and information kept on a datasheet describing the source of the blank. Results for all parameters should be non-detect. This type of blank is a check for cross contamination between sampling sites and control for bias introduced by cross contamination.

Other types of blanks will be used as needed: field sampling conditions or ambient blanks, if there is any reason to suspect that ambient air pollution has the potential to contaminate water quality samples; preservative blanks, if there is any reason to suspect that a preservative may be contaminated; or bottle blanks, any time sample collection bottles are of uncertain quality or cleanliness or from a source not previously used.

6.6.3.2 Sampling Duplicates

The purpose of a duplicate sample is to estimate the inherent variability of a procedure, technique, characteristic or contaminant. Duplicate samples are collected and duplicate analyses may be made in the field: 1) as a form of field quality control; 2) to measure or quantify the homogeneity of the sample, the stability and representativeness of a sample site, the sample collection method(s) and/or the technician's technique.

Duplicates are analyzed in the laboratory for the same parameters as the monitoring sample to which they apply. Laboratory duplicates which exceed QA/QC standards for the parameter are retested. Analytical results of duplicate samples will, theoretically, be the same. Realistically, results may differ due to the non-homogeneity of the sample source, and sampling and analytical errors. Duplicate samples also document the technique and ability of the technician and analyst to produce representative water quality data.

The laboratory analytical report must show test results for the duplicates, blanks and spikes, the method and the results for summary quality control statistics calculations. Copies of these reports are a permanent part of the site file.

Duplicates will ideally be splits of homogeneous samples to estimate measurement precision in the context of repeatability unless otherwise documented and justified. If enough sample water cannot be collected in the compositing container to facilitate a split sample, then duplicate samples will be co-located.

Duplicate field samples must be collected every sampling trip for each type of sample collected and the results must have a Relative Percent Difference (RPD) less than or equal to the guidelines in Table 6-6. Required field parameter measurements can be duplicated to estimate the precision of the equipment. Every tenth measurement may be duplicated, and the results of both measurements

recorded and evaluated as RPD. The result can be compared with the stated precision of the instrument.

Table 6-6. Frequency, acceptable range, and corrective actions for duplicate samples.

Type of Duplicate	Frequency	Acceptable Range for Precision	Corrective Action
Field duplicates (samples)	Minimum of 1 per trip per parameter or 10% of all samples per parameter per day	Chlorophyll-a, TSS and nutrients $\pm 30\%$ RPD; all other parameters $\pm 10\%$ RPD	Audit field personnel and verify sample collection procedure; resample; reanalyze; revise SOP; audit and train field personnel; project manager determines whether associated data is usable
Field duplicates (multi-probes)	Minimum of 1 per trip per parameter or 10% of all samples per parameter per day	All parameters $\pm 10\%$ RPD	Re-calibrate instrument; replace batteries; perform instrument field check with different standards; repair or replace instrument; notify management; audit and train field personnel; project manager determines whether associated data is usable

6.6.4 Summary of Quality Assurance/Quality Control Field Procedures

Quality assurance protocols are means to ensure data collected are as representative of the natural environment as possible. Quality assurance procedures are required in all data collection efforts as part of this monitoring protocol. Many of the key elements of quality assurance have been included in the SOPs where appropriate, as well as detailed in SOP #12. A summary of important QA/QC procedures follows. Table 6-7 lists additional requirements for QA/QC methods and documentation.

Table 6-7. Summary of QA/QC documentation and sampling methods.

Procedure	Description/reason
Instrument calibration logs	Each instrument must have a log in the form of a permanently bound logbook. Calibration schedule must be observed, using fresh calibration standards.
Project binder	Containing: checklist of QA/QC reminders, copies of decontamination, sample collection and processing SOPs, copies of equipment calibration and troubleshooting instructions, ASR and COC forms, blank field forms.
Site binders	Containing: GPS coordinates for verification of correct sampling location, table of previous field measurements to compare with new measurements
Field forms	Field forms are the only written record of field measurements, so copies are placed in site binders and originals must be kept on file indefinitely.
Field instrument methods	Require consistent measurement methods and detection limits
Sample preservation and minimum holding time	Water quality variable concentrations are maintained as close to sampling conditions as possible.

Table 6-7 (continued). Summary of QA/QC documentation and sampling methods.

Procedure	Description/reason
Chain-of-custody	A chain-of-custody includes not only the form, but all references to the sample in any form, document or log book which allow tracing the sample back to its collection, and documents the possession of the samples from the time they were collected until the sample analytical results are received.
Laboratory methods	Require consistent analytical methods and detection limits

- Field staff must be trained by personnel experienced in the protocol. Training will include classroom time as well as field and lab components. Details of the training are given in the Standard Operating Procedure #2: Training and Safety.
- Use calibrated instruments for all field measurements. Test and/or calibrate the instruments before leaving for the field. Each field instrument must have a permanent log book for recording calibrations and repairs. Review the log book before leaving for the field.
- All manually recorded field measurement data will be collected on field forms; data that are automatically recorded will be captured electronically and the equipment used will be documented on field forms. Hard and electronic copies will be made as soon as possible after surveys and kept at a separate location as backup.
- Complete records will be maintained for each sampling station and all supporting metadata will be recorded appropriately (field forms or electronically).
- Make field measurements in a manner that minimizes bias of results.
- Check field-measurement precision and accuracy. Follow the procedures in SOP #12.
- Collect 10% duplicate water samples; conduct duplicate measurements of field parameters at approximately 10%.
- Create field blanks prior to the beginning of each field season and periodically throughout the season.

6.6.5 Corrective Responsive Action

The project manager, in consultation with experienced professionals, will be responsible for taking corrective responsive action in the case in which QA/QC is not followed or in the case of an unexpected event. Responsive action is often needed in the event of broken sample bottles, missing data, errors on field sheets, changes due to field conditions, problematic analyses and other events that do not fall within the standard operating procedures. A “Memo-to-file” will be used to document any decisions or corrections that are made. This memo will include the date, name of author, site or sample referred to, a description of the problem or error and a statement describing the decision made or action taken. The memo will be archived with the appropriate site data and files.

6.7 Data and Records Management

- Hand-written field sheets with core suite profiles should be reviewed for completeness immediately upon returning to the shore. Make photocopies as soon possible and file them appropriately. Download and back up data from data loggers as soon as possible.
- Ensure that field forms, field notebooks, and other hardcopy records are secure, organized, and available for viewing, reproduction, or transfer upon request and/or at the end of each field season.
- Schedule and perform regular data transfer and backup. Data will be protected from loss or damage by daily backup, when possible, or on a feasible schedule approved by the project leader and the data manager. Ideally, raw data are backed up off-site as soon as a sampling trip is completed. Electronic data will be backed up from field sites using the GLKN FTP site following each sample round.
- Record and verify observed or measured data values. Complete paper forms and enter or import data into NPSTORET and/or other electronic databases.
- A ‘Memo-to-file’ will be used to document any decisions or corrections that are made. This memo will include the date, name of author, site or sample referred to, a description of the problem or error, and a statement describing the decision made or action taken. The memo will be archived with the appropriate site data and files.
- Data validation, the process by which data are proven or disproved to be accurate, involves the review of the results of all measurements, samples, and QC samples. Sample data should be flagged if the analyte was detected in a blank at a concentration similar to that in the sample; flagged data contain much more uncertainty than unflagged data. Some investigation into the sampling method may be needed if flagged data are a continual occurrence.
- Refer to SOP #8 for more details regarding data entry and management.

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SOP #6 Attachments

- Field Data Sheet
- Multiprobe Calibration Log
- Eureka Manta Maintenance Log
- Multiprobe Calibration/Maintenance Log

Park _____		Site Code _____		Date _____		Start time _____		End time _____		
Location Name _____				Secchi (m) 1st _____ 2nd _____		Ave. _____				
UTM/Lat/Long _____				Secchi reading cond.: excellent 1 moderate 2 poor 3 on bottom 4						
NAD _____		Zone*: 15N 16N WP# _____		Transparency Tube (cm) 1st _____ 2nd _____		Ave. _____				
Personnel _____				Water color (visual, in composite jug) _____						
Instrument(s) _____		Cal Date _____		Water sample type: I-tube Grab Van Dorn Other _____						
Model/Model# _____				VanDorn depth (m) _____		Wave Height (cm) _____				
Elevation (MSL, ft or m?) _____				Zmax (m) _____		probe other _____				
Barometric Pressure (_____ in Hg) (25.4mm/in) = _____ mm Hg				DO Cal (% sat.) Start _____		Finish _____				
B= Bottom, Q=QA/QC				Water Level Measurement						
Depth (m)	Temp (°C)	pH	DO (mg/L)	DO (% sat)	EC (µS/cm)	Initials	Observer 1: Benchmark Nail			
							Stadia Meas.			
							Correction (+/-)			
							Water Level 1			
							Initials	Observer 2: Benchmark Nail		
								Stadia Meas.		
								Correction (+/-)		
								Water Level 2		
							Avg WL 1 & 2			
							Water Sample Info			
Sample Volume Filtered (ml)										
Chl a _____ TSS _____										
Water Sample Collected for (check off):										
TP/TN, NOx/NH4 _____										
Chl-a _____										
Alk, Cl, SO4 _____										
DOC _____										
SiO2 _____										
(SACN only) Ca, Mg, Na, K _____										
(SACN only) TSS _____										
Other (diatoms, Hg, contaminants, etc.) _____										
Weather Observations - fill in blanks or circle code										
Current										
Wind Dir. (as °) _____		Wind Condition _____		Code _____		Past 48 hours				
Air Temp (°C) _____		Smoke rises vertically		0		Rainfall & Wind				
Sky Cover _____		Smoke drifts, leaves rustle		1						
Clear: < 10%		Wind felt on face		2						
Scattered: 10% to 50%		Flag extended, leaves move		3						
Broken: 51% to 90%		Dust, sm. branches move		4						
Overcast: > 90%		Small trees begin to sway		5						
Obscure: precip or obstructed		Large branches move, wires whistle, umbrellas hard to control		6		Any info on Back?				
Fog						Y N				
Light Mist		Beyond 6, you shouldn't be out in a boat								

[illegible][illegible]

Eureka Manta Multiprobe Calibration**Personnel****Date****DO Membrane Changed****Multiprobe****Date****pH Reference Electrolyte Changed****Time****Date**

	Calibration Standard (True Value)	Pre-Calibration Reading	Calibration Reading	Calibration Check	Pre- or Post-Field?
Battery Voltage					
Depth	0				
SpCond (µS/cm)	Air (0)				
SpCond (µS/cm)	value Lot#				
SpCond (µS/cm)	value Lot#				
SpCond (µS/cm)	value Lot#				
Temperature					
pH	7 Lot#				
pH	10 Lot#				
pH	4 Lot#				
pH	Lot#				
Baro Pressure					
DO% Saturation					
DO (mg/L)					

Eureka Manta Maintenance Log

Record maintenance performed, date, and person conducting maintenance

Date & Personnel	DO Probe	pH probe	EC25 probe	Temperature	Depth Sensor

MULTIPROBE CALIBRATION/MAINTENANCE LOG

PRE FIELD CALIBRATION					
Calibration _____ Initials: _____ Sonde ID: _____ Date: _____ Time: _____ Instrument: _____ Battery Voltage: _____					
Function	Temp. of Standard	Value of Standard	Initial Reading	Calibrated to	Comments
Specific conductance					
pH calibrated (~7)					
pH secondary standard (4 or 10)					
Dissolved oxygen					
DATA NEEDED FOR DISSOLVED OXYGEN CALIBRATION					
Altitude (A)= _____ feet above msl		Barometric pressure _____ inches			
Barometric Pressure (BP) Options		Barometric Pressure Formulas			
Barometer		Barometric pressure (inches) _____ x 25.4 = BP _____ mm			
From local source after correction (CBP)		BP _____ mm = CBP _____ mm - 2.5 (altitude ____/100)			
Estimated from altitude only		BP _____ mm = 760 mm - 2.5 (altitude ____/100)			
For older Hydrolabs: Table DO value _____ x ALTCORR _____ x BAROCORR _____ = DO standard _____					
POST FIELD CALIBRATION CHECK Post Calibration Initials: _____ Date: _____ Time: _____ Instrument: _____ Battery Voltage: _____ date of original calibration _____					
Function	Temp. of Standard	Value of Standard	Initial Reading	Calibrated to	Comments
Specific conductance					
pH 7					
pH secondary standard (4 or 10)					
Dissolved oxygen					
Check previous maintenance and use; do the following before calibration:					
Polish conductivity electrodes. Must be polished within the last two months or once every 15 field trips			Date:	Name/comments:	
Change pH reference probe solution. Must be renewed within last two months or once every 15 field trips.			Date:	Name/comments:	
Inspect DO membrane for nicks or bubbles. Must be changed within last six months or once every 15 field trips.			Date:	Name/comments:	
Change battery. Change once a year. Change internal batteries for newer generation products according to guidelines in product manual.			Date:	Name/comments:	

Standard Operating Procedure #7: Samples and Analytical Laboratory Requirements

Version 1.2

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Revision History Log

The following table lists all edits and amendments to this document since the original publication date. Information entered in the log must be complete and concise. Users of this standard operating procedure will promptly notify the project manager and/or the Great Lakes Network (GLKN) data manager about recommended and required changes. The project manager must review and incorporate all changes, complete the revision history log, and change the date and version number on the title page and in the header of the document file. For complete instructions, please refer to Revising the Protocol, SOP #13.

Previous Version Number*	Revision Date	Author (with title and affiliation)	Location in Document and Concise Description of Revision	Reason for Change	New Version Number
1.0	2/25/2010	Rick Damstra, David VanderMeulen, Joan Elias, GLKN	7.2 Sample Handling and Processing – clarification; 7.3 Shipping Procedures – clarification; 7.4 Quality Assurance/Quality Control - clarification	To bring SOP into alignment with procedures practiced	1.1
1.1	2/02/2016	David VanderMeulen, Rick Damstra GLKN	Updated NPS contact author and authorship, added info about TSS in tables; added that field samples might be collected as elbow deep grab samples, included information about acid-washing bottles, fixed a handful of grammatical errors, and other minor clarifications throughout.	To clarify and update SOP based on changes that have occurred since Version 1.1 was published in 2010	1.2

*Add rows as needed for each change or set of changes tied to an updated version number

Acknowledgements

Combinations of existing protocols were used to develop this standard operating procedure for handling and processing water samples prior to sending them to an analytical laboratory. The authors wish to acknowledge these sources: Hoffman et al. (2005), O’Ney (2005), USGS (2005), Water on the Web [WOW] (2004), Penoyer (2003), EPA (2001), and Baker et al. (1997) for the guidance they provided. Thanks also go to White Water Associates for providing examples of forms for analytical requests and chain of custody.

7.0 Introduction

As part of our water quality monitoring program, we will collect water samples for analysis of basic limnological parameters that may respond to stressors such as atmospheric deposition, land use change, and recreational pressures. Because neither the GLKN nor the individual park units currently have certified laboratories and instrumentation for performing these analyses, water samples will need to be processed soon after collection for transport to a suitable laboratory. This will require filtering and preserving subsamples of water and filters according to methods specific to the analyte. This standard operating procedure (SOP) is designed to provide detailed instructions on the handling and processing of water samples prior to analysis by an analytical laboratory.

We will collect water in the field (SOP #6) for the analysis of the following parameters:

- Nutrients: total phosphorus (TP), total nitrogen (TN), nitrate+nitrite-N ($\text{NO}_3+\text{NO}_2\text{-N}$), and ammonium-N ($\text{NH}_4\text{-N}$)
- Total chlorophyll-*a*
- Major anions (SO_4^{-2} and Cl^-);
- Alkalinity
- Major cations (Mg^{+2} , Ca^{+2} , Na^+ , K^+)
- Dissolved silica (SiO_2)
- Dissolved organic carbon (DOC)
- Total Suspended Solids (TSS)

Water chemistry will be performed by one or more analytical laboratories that have demonstrated the ability to measure analytes at detection levels adequate to meet our needs. Preferably, the laboratories will be state- or federally-certified for performing the above water chemistry analyses in natural waters, or an academic research laboratory that can demonstrate quality assurance and quality control (QA/QC) procedures consistent with SOP #12 and current EPA procedures used as the basis for state certification of commercial environmental laboratories. The GLKN's preference is for laboratories to provide clean sample bottles and preservatives, where appropriate, as well as chain of custody documentation and sample logging.

7.1 Summary of Analytical Methods

Analytical methods, method detection limits, and procedures related to handling samples change over time as the science progresses. Table 7-1 summarizes examples of the variety of methods, detection limits, preservation techniques, and holding times for water samples addressed in this protocol. Table 7-2 summarizes the frequency at which water samples will be collected for analysis.

Table 7-1. Examples of analytical methods, method detection limits (MDLs), containers, and holding times.

Analyte	Analytical Note 1	Method #	Det. Limit	Vol. (ml)	Filter	Preservation	Sample Bottle Note 2	Hold Time
Alkalinity	Titrimetry	310.1 EPA-NERL	10 mg/L	--	--	4°C	--	14 d
	Spec. auto.	310.2 EPA-NERL	10 mg/L	--	--	4°C	--	14 d
	Titrimetry	NFM USGS-OWQ	0.01 meg/L	--	Note 4	None	--	none
Ca ⁺²	ICP	3120B APHA	10 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
	Titrimetry	215.2 EPA-NERL	0.5 mg/L	--	Note 3	4°C	--	6 mo
	FAA	I-3152 USGS-NWQL	0.1 mg/L	250 mL	Note 3	pH<2 HNO ₃	P	180 d
Cl ⁻	IC	300.0 EPA-NERL	0.02 mg/L	--		4°C	P or G	28 d
	Colorimetry	325.2 EPA-NERL	1 mg/L	--		4°C		28 d
	Titrimetry	4500-Cl APHA	0.15 mg/L	100 mL		4°C	P or G	28 d
Chlorophyll-a	Spect.	10200 APHA	2 ug/L	≤1 L	Note 4	Freeze filter	P	30 d
Color	Spect.	110.2 EPA-NERL	5 Pt units		Note 5	4°C	G	48 hr
	Vis. Comp.	I-1250 USGS-NWQL	1 Pt-co	250 mL	Note 5	4 °C	P	30 d
DOC	Spect.	415.3 EPA	0.018 mg/L	125	Note 3	pH<4 H ₂ SO ₄	G	28 d
	Spect.	0-1122-92 USGS	0.1 mg/L	--		4°C	AG	
K ⁺	ICP	3120B APHA	0.3 mg/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
	FAA	3111B APHA	5 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
Mg ⁺²	ICP	3120B APHA	20 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
	FAA	3111B APHA	0.5 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo

Source: National Environmental Methods Inventory website (NEMI 2006)

This list is not an endorsement of any particular method or laboratory for any particular analyte. Rather it is to be used as a reference for the range of analytical methods available for each analyte. There are surface water conditions (pH, turbidity, other elements) that make a particular method unsuitable for a particular situation. As GLKN is monitoring surface water, the methods listed were chosen as representative of the lower range of detection limits.

Note 1. CIE-UV= capillary ion electrophoresis with UV detection, FAA = flame atomic absorption, FIA = flow injection analysis, IC= ion chromatography, ICP = inductively coupled plasma, Spec. auto = spectroscopy with autoanalyzer

Note 2. P = plastic (polypropylene), G=glass, AG=amber glass

Note 3. 0.45µm membrane filter. Pre-filter for dissolved portion analysis.

Note 4. 0.45µm glass fiber filter.

Note 5. 0.45µm membrane filter or centrifuge is recommended to remove suspended solids that affect color; however some color will also be removed.

Note 6. USGS 2003. Evaluation of Alkaline Persulfate Digestion as an Alternative to Kjeldahl Digestion for Determination of Total and Dissolved Nitrogen and Phosphorus in Water By Charles J. Patton and Jennifer R. Kryskalla. U.S. Geological Survey Water-Resources Investigations Report 03-4174.

Table 7-1 (continued). Examples of analytical methods, method detection limits (MDLs), containers, and holding times.

Analyte	Analytical Note 1	Method #	Det. Limit	Vol. (ml)	Filter	Preservation	Sample Bottle Note 2	Hold Time
Na ⁺	ICP	3120B APHA	30 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
	FAA	3111B APHA	2 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
NH ₄ -N	Selective elec.	4500-NH ₃ E	0.08 mg/L	--	--	4°C/pH2,0°C	--	24 hr / 28 d
	Colorimetry	350.2 EPA-NERL	0.08 mg/L	--	--	pH<4 H ₂ SO ₄	--	28 d
	Titrimetry	4500-NH ₃ APHA	5 mg/L	--	--	4°C / pH2,0°C	--	24 hr / 28 d
SiO ₂	ICP	3120B APHA	20 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
	Spect.	4500- SiO ₂ D APHA	0.04 mg/L	--	Note 3	No, 4°C	P	28 d
	FIA-Spect.	4500- SiO ₂ F APHA	0.78 ug/L	--	Note 3	No, 4°C	P	28 d
TSS	Gravimetric	I-3765 USGS-NWQL	1 mg/L	250– 500	Note 4	4°C filter	P	NA
SO ₄ ⁻²	IC	4110C APHA	75 ug/L	--	Note 3	pH<4 H ₂ SO ₄	P or G	--
	CIE-UV	D6508 ASTM	0.1 mg/L	--	Note 3	pH<4 H ₂ SO ₄	--	ASAP
	Spect.	37512 EPA-NERL	0.5 mg/L	--	Note 3	pH<4 H ₂ SO ₄	P or G	28 d
TP	Spect.	I-2606 USGS-NWQL	0.001 mg/L	125 mL		MgCl 4°C	BrownP	30 d
	Alkaline P	USGS 2003	0.01 mg/L	120 ml	Note 6	4°C / H ₂ SO ₄	--	48 hr / 30 d
	ICP	200.7 EPA-NERL	60 ug/L	--		pH<2 HNO ₃	P	6 mo
TN	Alkaline P	USGS 2003	0.03 mg/L	120 ml	Note 6	4°C / H ₂ SO ₄	--	48 hr / 30 d
	Titrimetry	4500-N	0–100 mg/L	--		4°C	AG	7 d
	Combustion	440.0 EPA-NERL	0.1 mg/L	--		Filter	--	100 d

Source: National Environmental Methods Inventory website (NEMI 2006)

This list is not an endorsement of any particular method or laboratory for any particular analyte. Rather it is to be used as a reference for the range of analytical methods available for each analyte. There are surface water conditions (pH, turbidity, other elements) that make a particular method unsuitable for a particular situation. As GLKN is monitoring surface water, the methods listed were chosen as representative of the lower range of detection limits.

Note 1. CIE-UV= capillary ion electrophoresis with UV detection, FAA = flame atomic absorption, FIA = flow injection analysis, IC= ion chromatography, ICP = inductively coupled plasma, Spec. auto = spectroscopy with autoanalyzer

Note 2. P = plastic (polypropylene), G=glass, AG=amber glass

Note 3. 0.45µm membrane filter. Pre-filter for dissolved portion analysis.

Note 4. 0.45µm glass fiber filter.

Note 5. 0.45µm membrane filter or centrifuge is recommended to remove suspended solids that affect color; however some color will also be removed.

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Table 7-2. Frequency of analyses of water samples and depths at which samples will be collected. Surface samples will be a 0–2 m integrated sample or an elbow-deep (ca. 0.3 m) grab sample, depending on site. Bottom samples will be collected during mid-summer if a site is thermally stratified.

Water Quality Variables	Monitoring Frequency
Total phosphorus (TP)	8x/yr, surface 1x/yr, bottom
Total Nitrogen (TN)	8x/yr, surface
Nitrate + Nitrite-Nitrogen (NO ₃ +NO ₂ -N)	8x/yr, surface
Ammonium-Nitrogen (NH ₄ -N)	8x/yr, surface
Total Suspended Solids (TSS)	8x/yr, surface
Major Anions (SO ₄ -2, Cl-)	3x/yr; surface
Alkalinity	3x/yr; surface
Major Cations (Mg+2, Ca+2, Na+, K+)	3x/yr; surface
Dissolved silica (SiO ₂)	3x/yr; surface
Dissolved organic carbon (DOC)	3x/yr; surface
Total chlorophyll-a	3x/yr; surface

7.2 Sample Handling and Processing Procedures

The following general techniques will be observed throughout the procedures detailed in 7.2.1 through 7.2.4.

1. Keep all water samples cool and dark until processing is complete and samples are shipped to the analytical laboratory.
2. Use only new, clean sample bottles supplied by the analytical laboratory, purchased pre-cleaned from a supplier, or those cleaned and acid-washed with 0.1N HCl by GLKN staff.
3. Rinse filtration equipment with deionized water (DIW) three times between samples.
4. Avoid touching the inside of sample bottles and filtering apparatus, tips of forceps, and filters to prevent contamination of the samples.
5. When filtering samples in the field, use an enclosed filtering apparatus to minimize contamination from airborne sources.
6. Wear disposable, powderless gloves when working with acids and other preservatives.
7. Filter samples in the order of anticipated phosphorus concentrations, from low to high. After filtering a water sample that is expected to contain high nutrient concentrations, rinse the apparatus three times with 0.1N HCl followed by three times with DIW water before processing the next sample.

8. Prepare QA/QC samples in the same manner as regular samples, using water from the same sample collection container.
9. Rinse all reusable equipment with DIW immediately, before equipment dries.
10. Ensure all sample bottles are labeled correctly, completely, and legibly.
11. Check laboratory equipment and supplies list (Table 7-3) and ensure equipment is clean and ready for use and supplies are adequate.
12. Prepare a temperature check bottle for each anticipated cooler, if recommended by the contract analytical laboratory. Use tap water to fill an extra bottle of the same size used for one of the analytes and label as “Temperature Check.” Store this check bottle in refrigerator with other samples; package and send to the analytical laboratory with the other samples.

Table 7-3. Laboratory equipment and supplies list.

• Filtration towers and manifold (4.7 mm) plastic
• Vacuum pump with pressure gauge and extra filtering flask as a water trap to protect the pump in case of overflow
• Graduated cylinders, plastic 250, 500 and 1000 mL
• Whatman GF/C filters (4.7 cm diameter)
• 0.45 µm Millipore membrane filters (4.7 cm diameter)
• Dried and pre-weighed TSS filter in appropriate container (Whatman GF/C filters, 4.7 cm diameter; provided by contract lab)
• Filter forceps with broad tips
• Aluminum foil
• Labeling tape, permanent markers
• Deionized water (ASTM grade 1 or 2; 1–10 megohm)
• Acid for preservation (according to contract laboratory specifications)
• Freezer
• Plastic storage bags
• Sample bottles (provided by analytical laboratory)
• Insulated ice chest, ice, and ice packs
• Saturated MgCO ₃ solution (for chlorophyll-a), depending on laboratory method
• Adjustable automatic pipettes: 1–5 mL ; 0.2–1 mL; 0.02–0.1 mL
• Paraffin paper roll
• Wash (squirt) bottles – 500 mL
• Kim wipes

The following sections detail the procedures to be followed when processing water samples for particular analysis.

7.2.1 Total Chlorophyll-a

1. Fit rinsed filtering device with a Whatman GF/C glass fiber filter using forceps, smooth side down (curl is up).
2. Agitate water sample (always shake well to minimize subsampling error for solids).
3. Set pump vacuum to ≤ 0.5 atmospheres (7.5 PSI or 380 mm Hg). If using a hand pump, maintain pressure at or below 10 PSI.
4. Use a glass or plastic graduated cylinder to measure 100–1000 mL of water sample. Filter sample. If water is very turbid, filter small aliquots (100 mL) to avoid clogging the filter. Sufficient volume has been filtered when a green, brown, or tan color is clearly visible on the filter and the flow decreases to a few drops/second.
5. Add 0.15 mL (approx. 3 drops) of saturated MgCO_3 during the last 30 mL of filtering to buffer the filter, if required by the method used by the contract analytical laboratory.
6. Rinse graduated cylinder and filtering apparatus with DIW and pass through filter to include any algae that may have adhered to the sides of the cylinder.
7. Record volume filtered on data sheet (excluding DIW rinse).
8. Use forceps to fold filter into quarters with sample on the inside; do not touch filter with fingers.
9. Wrap filter in foil; label foil with sample location, date, and time sample was collected, and volume filtered. Place foil in small, sealable baggie with standard laboratory label.
10. Refrigerate immediately and freeze as soon as possible. Place small baggies with foils together in a large, sealable freezer bag. A third watertight container may be used for shipping to ensure that melt water in transport will not corrupt the samples.

7.2.2 Total Suspended Solids

1. Use forceps to fit rinsed filtering device with a pre-weighed glass fiber filter supplied by the contract analytical laboratory.
2. Agitate water sample (always shake well to minimize subsampling error for solids).
3. Use a glass or plastic graduated cylinder to measure 100 - 2500 mLs of water sample. Filter sample. If water is very turbid, filter small aliquots (100 mLs) to avoid clogging the filter. Sufficient volume has been filtered when the flow decreases to a few drops/second.
4. Rinse graduated cylinder and filtering apparatus with DIW and pass through filter to include any sediment that may have adhered to the sides of the cylinder.
5. Record volume filtered on data sheet (excluding DIW rinse).
6. Place filter into container supplied by laboratory; do not touch filter with fingers.
7. Label container with sample location, date and time sample was collected, and volume filtered.
8. Refrigerate immediately. Samples may be stored refrigerated over the short term (1-2 months) or frozen if stored longer. Consult with the contract lab on their storage requirements. Place small baggies together in a large, sealable freezer bag. The large freezer bag may be placed in a watertight container for shipping to ensure that melt water in transport

will not corrupt the samples.

7.2.3 Unfiltered (Raw) Samples

1. Rinse sample bottle provided by analytical laboratory 1x with sample water.
2. Fill sample bottle with sample water (fill to just below neck if sample will be frozen).
3. Refrigerate or freeze, as per laboratory instructions, until packaging for transport to analytical laboratory.

7.2.4 Filtered Samples

1. Using clean forceps, place a 0.45µm Millipore cellulose membrane filter in the filtration apparatus. Rinse with 100 mL DIW into a cleaned (0.1N HCl and DIW rinsed as per sample bottle cleaning) filtering flask (glass or plastic). Rinse flask with filtrate and discard filtrate.
2. Filter a small amount (approx. 50 mL) of sample water; rinse filtering flask with filtrate and discard filtrate.
3. Filter enough of the sample to produce the required amount of filtrate to be tested.
4. Dispense the filtrate into separate bottles provided by the analytical laboratory as follows:
 - Unpreserved constituents—rinse bottle with small amount of filtrate (approx.. 10 mL) and discard; fill bottle to neck, refrigerate or freeze as per laboratory instructions.
 - Preserved constituents—if the sample bottle already contains preservative supplied by the analytical laboratory, fill bottle, and store at room temperature or refrigerate. If bottle does not contain preservative, rinse bottle with small of amount of filtrate (approx. 10 mL) and discard, fill bottle approximately $\frac{3}{4}$ full, add the preservative (ampule provided by laboratory) and continue to fill bottle until full. Gently roll bottle to mix. Store at room temperature or refrigerate.

7.3 Shipping Procedures

1. Call FedEx or other courier service ahead of time to arrange pick-up.
2. Make large quantities of ice cubes and ice blocks (or buy ice) ahead of time.
3. Line cooler with large plastic garbage bag.
4. Place all total chlorophyll-*a* baggies containing aluminum foil wrappers in one large sealable plastic bag. Place this baggie between 2 ice packs or bags of ice. It is critical that melt water does not soak the filters, so you may want to place large sealed bag of foil wrappers in a sealed plastic jar before surrounding with ice.
5. Use ice cubes, doubly bagged in plastic bags, to pack around samples; use other ice blocks (water bottles, soda bottles, etc.) as they will fit.
6. Include a temperature check bottle with the sample bottles in each cooler, if used by the contract analytical laboratory.

7. Complete the chain of custody (COC) form (example provided in Attachment A), keeping the “client copy” for the project files. Seal the laboratory’s copies in a one-gallon plastic sealable bag and tape to the inside cover of the cooler. Prepare separate COC forms for each cooler.
8. If the refrigerated samples are sent in the same insulated cooler with the frozen samples, protect them from freezing by wrapping them in newspaper, bubble wrap, or something similar.
9. Ship samples overnight so they are received the following day during a work-week, whenever possible. Contact the laboratory about Saturday shipment receipt availability before shipping samples on a Friday. Many laboratories do not have sample receipt staff on Saturday or charge extra for staff time.
10. Alert the contract laboratory when samples have been shipped via phone or e-mail. Be sure to get acknowledgement from the lab that they know the samples are en route. Ask the lab to let you know when the samples are received.

7.4 Quality Assurance/Quality Control

Quality assurance/quality control refers to all those things good investigators do to ensure their measurements are accurate (the absolute true value) and precise (reproducible, consistent), and that they include a good estimate of their uncertainty. It specifically involves following established rules in the field and laboratory to assure everyone that the sample is representative of the site, free from outside contamination, and that it has been analyzed following standard QA/QC methods. These methods typically involve comparing the sample to a set of known samples for estimating accuracy and replicating the measurement to estimate its precision.

In the context of field sampling and processing, quality assurance protocols are meant to ensure that data collected are as representative of the site as possible. Such procedures include implementing good field procedures and quality-control checks, careful post-collection processing of water samples to minimize artifacts due to contamination or mislabeling, proper storage and preservation techniques while in transit to a laboratory, and proper QA/QC by the laboratory itself. See SOP #12, Quality Assurance/Quality Control, for more details.

Quality assurance at the batch level (within the laboratory) is accomplished by using proper techniques and replicating 10% of the field samples.

The most important aspects of quality control in the collection of water quality samples are: 1) samples collected should represent the site at the time the samples are collected, such that the samples produce the quality of information necessary to meet the objectives of the survey; and 2) the integrity of the samples collected is not compromised by contamination, misidentification, or improper sample handling or preservation.

To help meet these quality control aspects, the transport and tracking of the samples from the field to the analytical laboratory that performs the chemistry analyses is critical. Each set of samples should include a Chain of Custody Form (COC), and if the lab uses it, an Analytical Services Request (ASR)

(see Appendix B for an example). To ensure correct processing of samples, the information recorded on the COC and ASR forms must correspond to each sample in the shipment. Check with individual analytical laboratory requirements for correct labeling codes and procedures. To prevent water damage to paperwork accompanying samples to the laboratory, place all paperwork inside a sealable plastic bag and tape the bag to the underside of the cooler lid. Keep a copy of the completed COC and ASR forms in the office binder.

7.5 Data and Records Management

Complete and accurate record keeping of field-derived data is an essential component of monitoring water quality. Field technicians, crew leaders, and project leaders share responsibility for collecting, verifying, and documenting data according to the guidelines in this monitoring protocol and all applicable standard operating procedures. Data and records management include the following responsibilities:

1. Refer to the GLKN Data Management Plan (Hart and Gafvert 2006) for overall guidance.
2. Follow the QA/QC procedures in SOP #8 for specific instructions on data entry, management, verification, and validation.
3. Record and verify observed or measured data values, including completing paper forms and entering data into NPSTORET and/or other electronic databases. NPSTORET maintains the necessary relationships between data values, equipment configuration and calibration, procedures, methods, and metadata.
4. Schedule and perform regular data transfer and backup. Data will be protected from loss or damage by daily backup when possible, or on a feasible schedule approved by the project leader and the data manager.
5. Review, verify, and correct field data and sample processing information as soon as possible after the actual survey (see SOP #8 for more details).
6. Prepare data and procedural documentation, especially deviations from the protocol or study plan, including metadata forms in NPSTORET and additional documentation requested by the project manager or data manager.
7. Ensure that field forms, field notebooks, and other hardcopy records are secure, organized, and available for viewing, reproduction, or transfer upon request.

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Attachment 7A. Example of a chain of custody.

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Water Quality Monitoring Protocol for Large Rivers, Version 1.1

Standard Operating Procedure #8: Data Entry and Management

Version 1.2

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Revision History Log

The following table lists all edits and amendments to this document since the original publication date. Information entered in the log must be complete and concise. Users of this standard operating procedure will promptly notify the project manager and/or the Great Lakes Network (GLKN) data manager about recommended and required changes. The project manager must review and incorporate all changes, complete the revision history log, and change the date and version number on the title page and in the header of the document file. For complete instructions, please refer to Revising the Protocol, SOP #13.

Previous Version Number*	Revision Date	Author (with title and affiliation)	Location in Document and Concise Description of Revision	Reason for Change	New Version Number
1.0	2/25/2010	Richard Damstra and David VanderMeulen, GLKN	8.3.2 Data Entry, Verification, and Documentation – update data checking procedures	To bring SOP into alignment with practices	1.1
1.1	1/05/2016	Richard Damstra, David VanderMeulen, GLKN	Updated the NPS contact author and authorship, distinguished discrete from continuous data, noted that EQUIS database is used to load data into the National STORET Data Warehouse, deleted outdated information related to data distribution and interaction with data, added a schematic to show data flow for continuous data, added information about the Integrated Resource Management Applications (IRMA) web portal, fixed a handful of grammatical errors throughout.	To clarify and update SOP based on changes that have occurred since Version 1.1 was published in 2010	1.2

*Add rows as needed for each change or set of changes tied to an updated version number

Acknowledgements

Primarily, we relied on the Great Lakes Network's data management plan in writing this document. Standard operating procedures for the entry and management of data written by other networks also provided guidance.

8.0 Introduction

Water quality data collected under the large rivers water quality monitoring protocol must be entered, quality-checked, documented, managed, and made available to others for a variety of purposes, such as management decision-making, research, and education. The National Park Service (NPS) Water Resources Division (WRD) developed a database application called the Water Quality Database Template, or NPSTORET, to facilitate management of water quality data. NPSTORET facilitates the importing of water quality data into a copy of Earthsoft's EQuIS database maintained by WRD. From the EQuIS water quality data is imported into the National STORET (STORage and RETrieval) Data Warehouse administered by U.S. Environmental Protection Agency (EPA), as required by the water quality component of the Natural Resource Challenge (NPS 1999), NPS-75, and NPS Reference Manual 77. NPSTORET is best suited for water quality data that comes from discrete sampling during individual site visits.

To better accommodate continuous or time-series data that are generated from automated data loggers, WRD adopted Aquatic Informatics' Aquarius system for storage and visualization of continuously monitored water quality data. The Network will utilize a version of Aquarius hosted by WRD for remote data entry, analysis and visualization.

This standard operating procedure outlines data stewardship roles and responsibilities, and provides specific instructions and references for entering, quality checking, and managing water quality data.

8.1 Data Stewardship Roles and Responsibilities

Data stewardship is for the process of managing data and information resources that are organized, useful, compliant, available, and safe. The demand for detailed, high quality data and information about water quality requires a group of people working together to ensure that data are collected using appropriate methods, and that resulting datasets, reports, maps, and other derived products are well-managed.

The Great Lakes Network (GLKN) aquatic ecologist serves as project manager for water quality monitoring. The project manager will supervise data collection, provide project oversight, direct on-the-ground data collections, and provide cohesive links among data collection, synthesis, interpretation, and reporting.

While the project manager must act as the steward for water quality monitoring data for the Network, other project and GLKN personnel are also accountable for specific data management tasks. Table 8-1 lists stewardship responsibilities of personnel involved in the management of water quality data. To ensure that all project data are managed properly, individuals must understand their responsibilities, communicate with one another, and assist one another as needed.

Table 8-1. Data stewardship responsibilities of water quality monitoring personnel.

Personnel Role	Data management responsibilities related to water quality monitoring
Project Crew Member	Crew members collect and manage data with direction and guidance from the crew leader and/or project manager. Data collection includes calibrating and operating sampling equipment, collecting water samples, and recording measurements and observations. Crew members are responsible for quality control by following data collection and recording instructions and by promptly verifying recorded data. Crew members may also perform data entry and verification.
Project Crew Leader	The crew leader normally performs the same duties as the other crew members and ensures adherence to data collection and processing protocols, including data verification and documentation. The crew leader also works with the project manager and data manager on water quality data management in the office.
Project Manager (Network Aquatic Ecologist)	The project manager is responsible for all project operations and results, and may also participate in field operations. The project manager ensures that data management activities are conducted according to established procedures and is responsible for data validation and certification: approving the data content, quality, and documentation, as well as making decisions about data sensitivity and distribution. The project manager is responsible for evaluating project data at specified intervals, analyzing data for trends, and following reporting requirements.
Resource Specialist	The water resource specialist, who may also serve as a crew leader, works closely with the project manager in all aspects of data management. The water resource specialist collects field data, enters field and laboratory data, verifies data, and validates data.
Network Data Manager	The network data manager, together with the project manager, ensures that water quality monitoring data are organized, useful, compliant, available, and safe. The network data manager provides the most current version of NPSTORET and works with project personnel to ensure Network water quality data are received by WRD. The network data manager oversees activities related to training, user support, quality assurance, documentation, backups, archiving, and data maintenance and distribution.

8.2 NPSTORET Database

Water quality data are managed according to guidelines from the NPS Water Resources Division. In accordance with these guidelines, the desktop database application NPSTORET will be used to enter, store, document, and transfer discrete (or snapshot) water quality data. The GLKN oversees the use of NPSTORET per the Network's water quality monitoring protocols and ensures that data are transferred at least annually to the NPS Water Resource Division for upload to the EPA STORET database/Water Quality Exchange (WQX) (Figure 8-1).

STORET is an interagency water quality database developed and supported by the Environmental Protection Agency (EPA) to house local, state, and federal water quality data collected in support of managing the nation's water resources under the Clean Water Act. NPS Director's Order 77 indicates that the NPS should archive water quality data in STORET, and the NPS WRD mandates that any data collected as part of a WRD-funded project get archived in STORET. The NPS uses STORET as a repository of physical, chemical, biological, and other monitoring data collected in and around national park units by park staff, contractors, and cooperators. The NPS operates its own Service-wide copy of STORET and makes periodic uploads to the EPA STORET National Data Warehouse so that data collected by and for parks will be accessible to the public.

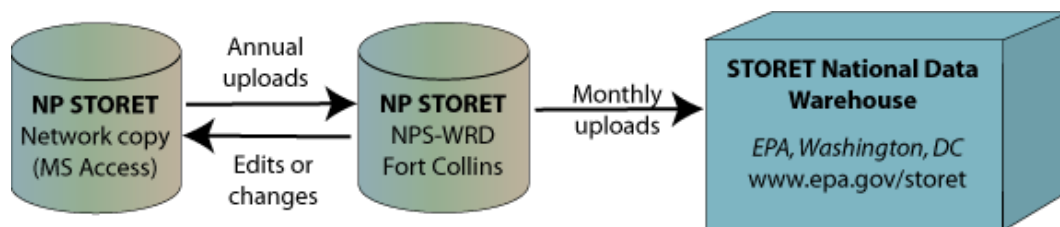


Figure 8-1. Schematic of data flow to the STORET database: water quality data are transferred from the Network to the national master database and ultimately to the master database. Public access to STORET is available via an online clearinghouse.

8.2.1 Database Design

The database design for NPSTORET is described in its associated documentation. Maintenance and documentation of NPSTORET is performed by NPS WRD. Because it must be compatible with the Oracle-based EPA STORET, NPSTORET is a complex MS Access implementation. The latest version of NPSTORET can be found at: <http://nrdata.nps.gov/programs/water/npstoret/>.

The Great Lakes Network will maintain one master copy of NPSTORET for each park at the Ashland office on a central server. This is the only copy of NPSTORET that can be used to export data to other locations (i.e., WRD and GLKN's SQL Server). Additional copies of NPSTORET can be used by GLKN personnel stationed at parks, but they can only be used as a conduit for data entry and the importation of data to GLKN's master version of NPSTORET. For analysis, the data from the master copy of NPSTORET, that has passed all QA/QC procedures, must be used.

8.3 Aquarius Database

Guidelines for managing time series data are being developed by NPS Water Resources and Inventory and Monitoring (IMD) divisions. For water quality monitoring personnel, data collection, management, and stewardship responsibilities for time series data are the same as those for discrete or snapshot data (Table 8-1). At the time of this latest SOP revision (Version 1.2), WRD and IMD are in the process of creating SOPs to use Aquarius tools to import, QA/QC, and validate and certify continuous data. A general data flow schematic for continuously collected data is shown in Figure 8-2.

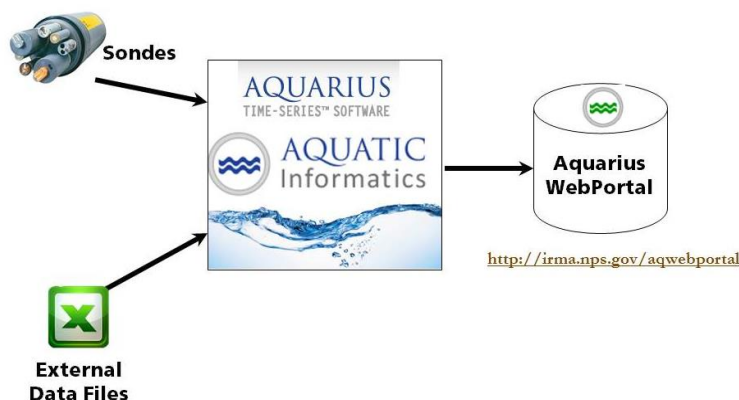


Figure 8-2. Schematic of data flow to the Aquarius database: water quality data from data loggers are transferred from the Network to the WRD-hosted version of Aquarius. Public access to this data is provided through the Aquarius WebPortal.

8.4 Data Management Procedures

8.4.1 Data Collection

Data values are measured, observed, or estimated according to the GLKN large rivers water quality monitoring protocol at various monitoring locations (sample sites) and recorded on field forms (see SOP #6, Field Measurements and Water Sample Collection). Crew members are responsible for legible, accurate entries on field forms and in log books, including the calibration log. As a first step to verify data, crew members will check and double-check the recorded values on the day of data collection.

Discrete data collected with a multiparameter sonde are stored directly on a datalogger attached to the sonde and recorded on the field sheets at the time of sampling. The hard copy of the data serves as a back-up should something happen to the electronic data.

Continuous data collected by temperature sensors through use of thermal arrays are stored directly on the sensors, downloaded to a data transfer device (e.g., a Hobo Shuttle), and transferred to a computer. Hard copies of continuous temperature data do not exist, although electronic data are stored in multiple locations and backed-up daily.

Digital images of sample sites are acquired during site establishment and periodically as sites change (SOP #6). Crew members are responsible for proper settings and use of digital camera equipment and should refer to the user manual for details specific to the camera.

GPS coordinates are stored as waypoints if using a recreational GPS unit, or as features if using a mapping-grade unit with a data dictionary, and recorded on the field sheets. When possible, the GPS data will be differentially corrected to improve the accuracy of location coordinates. See SOP #3 for more information on using a GPS unit.

Water samples are collected, labeled, and packaged for laboratory analysis according to SOPs #6 and #7. Identification numbers on sample containers, chain of custody forms, laboratory reports, and on the field data collection form facilitate management of laboratory results.

8.4.2 Data Entry, Verification, and Documentation

Requirements for data entry into NPSTORET are detailed in the documentation for users. We present a summary of data entry below, and refer staff to the most recent NPSTORET documentation for the specifics.

On a regular schedule approved by the project manager and data manager, the crew leader gathers the field data collection forms and verifies the completeness, accuracy, and legibility of each form. Following each round of sampling, the crew leader will make a photocopy of each field form and ensure each copy is legible; the copy will be placed in the office binder kept at the park, and the original will be sent to the project manager for archival in the GLKN office. Additional photocopies will be made as needed. Upon receiving the original data sheets, the project manager will proofread the datasheets, making sure that they have been filled out completely. All data sheets should have been reviewed for completeness while in the field, however some deficiencies in data recording may not be identified until all data sheets have been reviewed as a group.

Project staff enters site establishment data in the NPSTORET Station Entry Template as soon as possible following the initial site visit to each sample location. This prompt action is a good data management practice and an NPSTORET requirement that enables parameter data entry. Linking digital images of sample sites to stations in NPSTORET is required as part of this data entry process.

File size for digital images linked to stations in NPSTORET should normally be at least 100kb and less than 300kb. Project staff will reduce the size of image files larger than 300kb and copy images to a file folder named “images” for the appropriate park. Original ‘raw’ images can be stored in the Images folder (see Folder and File Organization below) if project staff determine it is important to keep higher resolution images.

Electronic data from the multiparameter sonde are downloaded from the datalogger to a computer in an MS Excel spreadsheet or text file at the end of each sampling day or as soon as feasible after sampling, and verified for accuracy. Likewise, at the end of each sampling day or as soon as feasible after sampling, all digital images will be downloaded to a computer and labeled with date, location, and subject matter. Import data from the MS Excel spreadsheet into NPSTORET following the guidelines.

Any notes taken in the field regarding collection of data with the sonde are transcribed to NPSTORET in a field for comments and notes. Results of laboratory analyses typically are sent to the project manager in a MS Excel spreadsheet and on hard copy forms. The project manager will verify both sets of data and follow-up with the contract laboratory if any discrepancies are noted or any questionable results are reported (see SOP #12 on QA/QC for more details). Project staff will import the verified data into NPSTORET.

Non-electronic field data recorded on hard copy data sheets are manually-entered into a MS Excel spreadsheet. At the end of the season when all data are entered, the project leader will perform or coordinate a check of 100% of this manually-entered data prior to import into NPSTORET.

A random check of 3% of all electronically transferred data (GPS, multiprobe, and laboratory data) is also performed after it is imported into NPSTORET, to ensure no problems occurred during data transfer. For electronically transferred or manually-entered data, the project leader resolves errors according to established procedures. See SOP #12 for additional QA/QC details.

Data verification starts with the QA/QC steps that are detailed in SOP #12. As data are entered into NPSTORET, either by keyboard or using an import routine, a suite of QA/QC procedures exist that compare the entered data with expected formats and accepted data value ranges or domains. If the entered data do not pass a form-based QA/QC test, NPSTORET prompts the user to make corrections and re-enter the data. For most of the form-based tests, the NPSTORET database will not accept out-of-bounds data, and correction is mandatory; for some range tests, out-of bounds data are accepted after a user prompt, but are flagged accordingly. The QA/QC procedures on imported data perform similar tests, but exceptions that do not pass QA/QC are presented to the user and must be reconciled before the record or dataset can be accepted by NPSTORET.

Additional comparison tests will be run on data in NPSTORET to flag records that are outside the expected rate of change for a parameter based on previous records for that parameter. These suspect data points are reviewed by an expert user and can be corrected, flagged, or excluded from the dataset.

As mentioned above, NPSTORET performs numerous QA/QC checks on the data as they are entered and stored. Additionally, once the data are transferred to WRD, more QA/QC procedures are performed. The Network's water quality data records are regarded as being in provisional status until they are returned to GLKN from WRD, or are accepted by WRD without changes after the final QA/QC steps. Each individual record in the water quality dataset always has one, and only one, status flag indicating its status as provisional or final. Finalized data can still be edited and changed if errors are discovered after review in reporting and analysis, but the status of those records reverts to provisional and they must be resubmitted to the master STORET version at WRD and returned before they can be reassigned as finalized data records.

A user's identification is assigned to each aspect of data handling, from collection through the final steps of QA/QC. On data sheets, the identity of the data collector(s) is recorded and is transcribed into the data entry forms in NPSTORET. Additionally, NPSTORET records the user name (login) for every table entry or modification. Only qualified users who have been trained and given edit permissions are allowed to edit data in NPSTORET. These procedures protect the integrity of the data and allow the history of each data record to be traced.

The Aquarius database software used for continuous water quality data has built-in QA/QC tools specifically tailored for time series data. These features tools are particularly useful for dealing with data gaps, overlaps and out of range values, and will be utilized as part of the QA/QC process used

for continuous data imported into Aquarius. At the time of this SOP development WRD and IMD are working cooperatively to create SOPs for QA/QC and data certification (see next section) for the Aquarius tool. GLKN will adopt the SOP's for these tasks when they are completed.

Executive Order 12906, mandates federal agencies to "...document all new geospatial data it collects or produces, either directly or indirectly..." using the Federal Geographic Data Committee Content Standard for Digital Geospatial Metadata. Water quality monitoring meets the definition of geospatial data and, thus, GLKN is responsible for documenting all public datasets using metadata that meets that standard. NPSTORET and Aquarius require and store a great deal of metadata at the record and project level. Metadata will be developed, using NPSTORET and Aquarius as primary sources that meet the federal standard. Consult the GLKN Data Management plan (Hart and Gafvert 2006) for additional details about metadata procedures and requirements.

8.4.3 Data Validation and Certification

Prior to distributing the data for any type of use, the project manager or other appropriate water resource specialist validates the NPSTORET database content in the master water quality data file according to procedures in the Quality Control and Quality Assurance SOP. The integrity of each master data file must be preserved during the validation process in cases where the person performing data validation does not have direct access to the master water quality data file. Validation is performed at least once each year after data collection and entry are complete and before data are submitted to the NPS Water Resource Division, usually by the end of each calendar year. If this annual validation does not meet scheduled reporting requirements of the Network, then the project manager may coordinate more frequent validation to meet needs, or reports can include a statement explaining that results are based on data that have not been validated. The statement should include an explanation of what significance this has for using preliminary data.

Laboratory results must be imported and validated in NPSTORET as soon as possible following receipt of the results. The project manager coordinates receiving and entering data with project staff and then performs or coordinates the validation of lab results.

Data certification will be performed annually according to policies in the Network Data Management Plan, Hart and Gafvert (2006). The certification process will take place before the finalization of the annual NPStoret submission to WRD, and the release of data in Aquarius for public viewing.

8.4.4 Data Analysis and Reporting

Project staff will follow the procedures for data analysis in SOP #9 and data reporting in SOP #10. Data summary statistics will typically include, but not be limited to:

- Mean
- Median
- Standard deviation
- Minimum
- Maximum

- Count
- Percentiles (10, 25, 75, and 90th percentiles)
- Standard error
- Variance
- Range
- Mode
- Sum
- Kurtosis
- Skew

Some of these statistics are available through NPSTORET's query function and others can be derived from NPSTORET outputs. NPSTORET provides an export for any portion of the data in tabular format for use with the Data Analysis Toolpak.

Aquarius has primarily been a tool used by hydrologists to compile stage and discharge data for flowing waters. It can generate the same data summary statistics as NPSTORET as well as a few others, such as reports by location showing minimum, maximum, mean, and count number of data points per time series. Aquarius can also be used to calculate means over several time scales (monthly, daily, or hourly) and to generate time series graphs. In addition to these and other analysis procedures, data can also be exported from Aquarius to flat file formats for use by other graphing or statistical software, such as R or SigmaPlot.

8.4.5 Data Folder and File Organization

All data from this water quality protocol should be stored, at the earliest possibility, on the GLKN central server. A diagram showing the folder structure is shown in Figure 8-3.

Files should be named in accordance with the file-naming standards in the GLKN Data Management plan (see section 6.4 of Hart and Gafvert 2006). Files have a 'GLKN' prefix, a descriptive element, and finish with a date element. For example, 'GLKN_ISRO_Field_Data_20070605.doc' contains field data from Isle Royale National Park on June 5, 2007. Do not use spaces in file names.

8.4.6 Data Archival and Distribution Procedures

Data archiving serves two primary functions: it provides a source to retrieve a copy of any data set when the primary dataset is lost or destroyed, and it provides a data record that is an essential part of the QA/QC process. Original data will be archived at the Network office. Original data for printed forms are either the physical datasheets or exact and complete digital copies of the forms that capture all entries and notations. The unedited files are the original data for digital data.

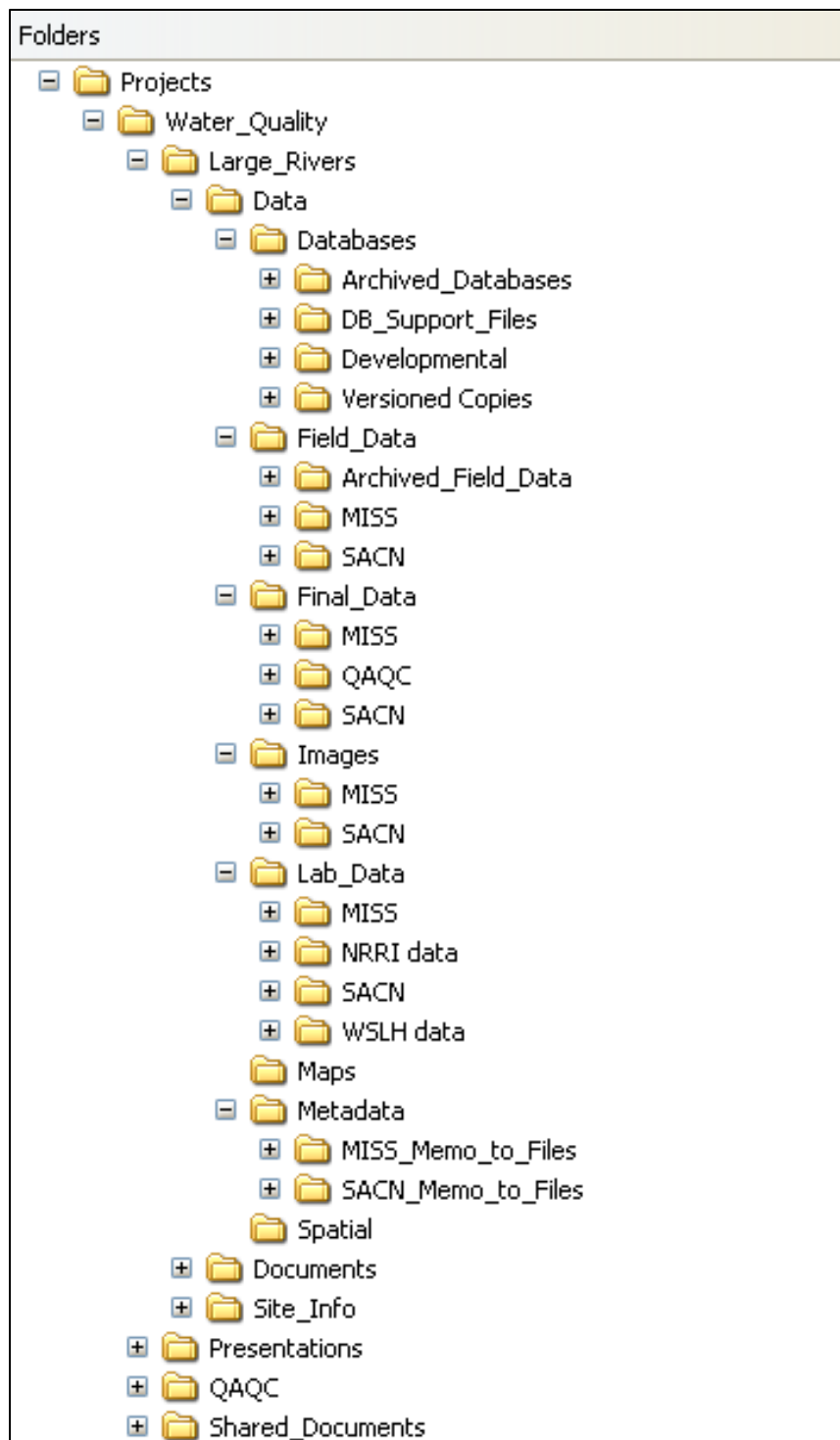


Figure 8-3. Great Lakes Network folder structure for information related to the protocol for monitoring water quality of large rivers. The Images, Lab_Data, and Maps folders also have subfolders for each park. Data are organized in subfolders by year within each park folder.

All digital data have a duplicate file created at the earliest opportunity. At least two complete copies of any water quality dataset are required by WRD, including digital replicas (scanned versions), if

they are created, of hard copy data sheets. Digital field data that are entered directly into a field computer or collected from a data logger must be backed up to a second medium at the earliest possibility. The data files on field computers and loggers must not be erased until the integrity of these data files are verified on the duplicate storage medium. The removal of original data files from a field computer or logger must be a balance of keeping memory available for new data collection and a need to keep data in their most original form for as long as possible. Field files should only be deleted when memory space is needed for new data collection.

The Network's master version of NPSTORET and the SQL Server geodatabase are maintained on a central server in the Ashland Office that is backed up daily, and backed up off-site weekly. Complete details of the GLKN Server archiving procedure are found the Infrastructure chapter of GLKN's Data Management Plan (Hart and Gafvert 2006); the general strategy for data archiving is also described in this plan and its appendices.

Raw time series data (.hobo files and flat files created from the .hobo files) are stored on the central server at GLKN, which is backed up daily, and backed up off-site weekly. The version of Aquarius that GLKN uses is housed on a server at WRD in Ft. Collins, Colorado. For the time series data to be stored and used in Aquarius, the flat file versions are uploaded to the SQL server using the Aquarius Springboard. All Aquarius data is uploaded to this server at least annually by GLKN staff or cooperators. WRD is currently working on a public-facing web application where this time series data will be available, once the data is officially certified.

GLKN has a number of different avenues for data distribution depending on the audience served and the degree of analysis and customization needed by the end-user (see GLKN's data management plan, Hart and Gafvert [2006], for more details). The NPS IMD Integrated Resource Management Applications (IRMA) web portal is the primary tool that provides interactive access to GLKN data. Some GLKN water quality datasets are available through links to the IRMA portal on the GLKN webpages. However, due to the complexity of the NPSTORET database, providing the data in that format will not meet the needs of those users unfamiliar with that application. Network (GLKN) versions of NPSTORET are used to update a master version of STORET maintained by NPS WRD. The WRD master copy of STORET data is used to regularly update the EPA database that provides on-line access to GLKN water quality data through the EPA WQX and forthcoming Water Quality Portal. NPS IMD is also developing a web map portal to monitoring data tentatively called the NPS Data Visualizer. This will allow a user to spatial query data, including monitoring data, and retrieve the scope of interest. The pathway for this tool is via a Common View Data Template that will allow data in NPStoret to be read by the database server supplying the Data Visualizer.

Data distribution will primarily occur through the EPA STORET portal, the NPS Data Store, Aquarius WebPortal and the NPS Data Visualizer. These tools are described previously in this document. The USGS and EPA are developing the Water Quality Portal (www.waterqualitydata.us). This will probably become the primary tool for reaching data stored in EPA STORET/WQX. For many users, the EPA tools will be the easiest way to retrieve subsets of GLKN large rivers water quality data.

In cases where specific data are requested that are unavailable through the public distribution methods outlined above, the project manager or appropriate water resource specialist will respond based on the nature of each request and the state of the data, for example, whether or not the data are fully qualified and documented to meet the needs of the requestor.

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Standard Operating Procedure #9: Data Analysis

Version 1.1

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Revision History Log

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Previous Version Number*	Revision Date	Author (with title and affiliation)	Location in Document and Concise Description of Revision	Reason for Change	New Version Number
1.0	3/17/2016	Richard Damstra, David VanderMeulen, GLKN	Updated the NPS contact author and authorship, distinguished discrete from continuous data, fixed a handful of grammatical errors throughout	To clarify and update SOP based on changes that have occurred since Version 1.0 was published in 2008	1.1

*Add rows as needed for each change or set of changes tied to an updated version number

Acknowledgements

We acknowledge Brenda Moraska Lafrancois and Richard Axler for helpful input and discussions relative to analysis of water quality data and general limnological principles. Thanks also to Erik Beever for providing language in earlier drafts of this document.

9.0 Introduction

Data analysis is the process by which measurements of the environment are interpreted meaningfully. It begins with evaluations of data after the data have been collected and entered into an electronic file or database, and have undergone a check of the data entry to ensure quality. Data analysis includes quality assurance/quality control (QA/QC) checks for statistical outliers prior to data summarization, may include exploratory data analysis, and concludes with analyses that lead to summary and interpretations of the data.

The appropriate method for analyzing monitoring data is tightly linked to the monitoring objectives, the spatial and temporal aspects of the sampling design, and the expected uses of the data by managers (MacCluskie et al. 2004). Methods of data analysis need to be considered well before the data are even collected.

As described in the GLKN protocol narrative for monitoring water quality in large rivers, the sample design of this protocol includes both targeted and probabilistic sampling. In other words, it includes some sites that are selected nonrandomly and others that are selected randomly. These two types of sites represent independent domains within the population of waterbodies that GLKN has defined as ‘large rivers’ within its nine park units. Therefore, each type of site will require its own method of data analysis (discussed below).

9.1 Initial QA/QC Checks for Outliers in Data

All water quality data undergo several quality assurance/quality control (QA/QC) procedures (e.g., duplicate sampling, data-entry filters, removal of logically inconsistent entries; see SOP #8 for data management details) to ensure that the data accurately represent the natural environment at the time of sampling. In addition to these procedures, several analytical and data sorting techniques (e.g., scatter plots, box-and-whisker plots, stem-and-leaf plots, sorting values of a given indicator in ascending or descending order, “COUNTIF” statements in Excel) are available to identify potentially erroneous values and statistical outliers. During this process, data points that do not meet limitations for precision or bias may be flagged or eliminated from the database. Some statistical measures, such as population mean, are sensitive to extreme atypical values, or outliers. Therefore, to reveal the central tendency of the population, the project manager may elect to remove outliers from the data pool. Such removals should be performed with great caution, and only when it is clear that the outlier truly did not reflect system properties (e.g., the outlier resulted from instrument error, transcriptional error, contamination of the sample).

9.2 Adjustment of Flow

Flow directly influences the values of most water-quality variables; many metrics are expressed in units of concentration (i.e., mass per unit volume). The purpose of monitoring water quality is often to detect change in the total mass of a variable, independent of volume of water present. Ideally, without budget constraints, this mass is directly calculated from continuous records of flow and high-frequency water sampling. However, in the face of budget constraints, flow-adjustment is another

method to account for the effect of flow. Flow-adjusted concentrations are thus not subject to the confounding influence that variable flow rates can produce. In particular, one uses regression or a smoothing technique (e.g., LOWESS) to obtain residuals from a relationship of concentration vs. flow (on a log-log scale). Trend analysis is subsequently performed on the residuals, which are presumably de-trended for flow. The trend in the residuals can be back-transformed to yield the trend in concentration units per year.

9.3 Spatial Domains of Analysis

Whereas some analyses may analyze trend at individual stations only (e.g., to investigate the effect of a local, point-source contamination), the most convincing conclusions will usually derive from results obtained across a collection of sites. Alternative domains of analysis include a portion of a park unit (e.g., the upper St. Croix River), an entire park unit, all monitoring stations on GLKN non-wadeable rivers, or regional or national domains. In these cases, individual stations constitute replicates, and the variability among stations provides an indication of the consistency in values across the larger landscape (within the chosen domain). Analysis of trend thus indicates whether a given water-quality parameter is changing or not *across an entire region*. Domains can be constructed to include several different sub-regions of, for example, a particular park unit. These sub-regions should be based on dominant physical or ecological gradients that are meaningful to the target indicator and not expected to vary through time. Thus, to the extent that stratification reduces the CV within parameters of interest, it will increase power to detect change over time, if total sample size is kept constant. It should be remembered, however, that if sampling units within a given domain are not selected probabilistically (with respect to how the domain is specifically defined), extrapolating results to broader domains should not be performed. Accordingly, appropriate caveats must be provided for interpretation of results provided.

9.4 Annual Data Summaries

Brief characterizations of the data from each park, and the Network as a whole will be performed annually, after all QA/QC procedures have been completed. Site-specific descriptive statistics may include mean, median, maximum and minimum values, as well as skew, kurtosis, and measures of variability, when appropriate (e.g., coefficient of variation, standard error, variance), and provided as hard copy or web based summary reports, with the primary audience being the parks

Given the relevant legislation (e.g., Clean Water Act of 1972), it may be of interest to individual NPS units and to other entities to assess the proportion of measurements during a time period or across a domain (at a single point in time) that exceed pre-determined thresholds, such as state water quality standards or ecoregional nutrient criteria. As with nearly all percentage data, arcsine transformations must be performed on those data before statistical analyses can be conducted (Sokal and Rohlf 1995).

9.5 Analyses of Long-term Trends

Following collection of at least three years of data at a given park, analyses more detailed than the data summaries described above may be conducted. Approaches to such detailed analyses may

include estimation of interannual change, graphic approaches (e.g., comparison of mean and variability in a parameter in the current year versus during past years), and occasionally qualitative analysis (Guthery et al. 2001), as well as modeling, correlational analyses, and various parametric and nonparametric analyses.

9.5.1 Recommended Methods for Long-Term Trend Analysis

After the first three sampling years of monitoring data are collected at Mississippi National River and Recreation Area (MISS) and St. Croix National Scenic Riverway (SACN) and periodically thereafter, more-intensive analyses of change will be performed. In addition to repeated-measures, time-series, regression, and non-parametric equivalents of various methods such as ANOVA and *t*-tests, monitoring data may also be evaluated through Monte Carlo simulation analyses, Bayesian analyses, and comparisons of period means. For the latter-most approach, one is often interested in comparing values before and after an important event (e.g., changes in management policy, remarkable anthropogenic disturbance, natural catastrophe, or drought). Years are then considered as replicates within each of the two periods. Trends in parameters will be analyzed with respect to biotic and abiotic covariates, although cause-effect relationships may be investigated more thoroughly by NPS partners and collaborators (e.g., USGS-WRD, university investigators).

In addition to analyzing each variable separately, flow-adjusted water-quality variables that are not correlated and that naturally could be considered a homogeneous group of parameters could be analyzed collectively through multivariate ordinations (e.g., nonmetric multi-dimensional scaling, NMDS; McCune and Grace 2002) of resource conditions through time. This approach effectively integrates information across many indicators, and can suggest whether individual stations are all moving in the same direction in multidimensional ordination space. Furthermore, joint plots can be overlaid on the ordination, and can suggest which variables correlate most strongly to the direction of changes. Similarly, if specific comparisons were desired across a suite of uncorrelated variables, two or more pre-defined groups of samples could be compared using MANOVA or NPMANOVA, depending on whether parametric assumptions are met (Sokal and Rohlf 1995; Zar 2007).

9.5.2 Approaches to Analyze Frequently-Collected Data

In contrast to many other national parks and the water quality monitoring at associated inland lakes, water quality measurements from the three non-wadeable rivers are (and have been for many years) collected many times per year. Not only does the long-term nature of this dataset allow for robust retrospective analyses of trend, but the multiple sampling sessions within each year allow for various approaches to analyze long-term trends. For example, if the goal is to monitor trend in the *average* value of a particular water-quality parameter, then all measurements collected within the year at each sampling location would be considered temporal subsamples and would be averaged. Trend analysis (using repeated-measures, time-series, or other regression analyses) would thus be performed simply on the annual means at each sampling location. This approach seems particularly viable for parameters that do not exhibit strong intra-annual variability (e.g., pH). In contrast, this approach may be vulnerable to imprecision or bias if a parameter exhibits significant, predictable variability in its value throughout the year and data are not collected at a high number of intended sampling occasions.

Alternatively, if a particular parameter is known to exhibit significant intra-annual variability (e.g., phosphorus concentration), samples within the year can be partitioned into one of several periods (ideally, defined by relevant phenological or biological phenomena). Thus, for example, if interannual trends in nutrient concentrations during algal blooms are of interest, the temporal domain can be accordingly defined, and all samples within that window averaged within each year and the means analyzed for trend across years. Finally, if there is a strong desire to incorporate intra-annual variability into interannual trend analyses, data within each year can be analyzed through a smoothing algorithm, and interannual variability is thus analyzed on the smoothed data. It may be the case, however, that process variation is larger than the sampling variability (Burnham et al. 1987).

9.5.3 Duplicate Sampling

To ensure that understanding of water quality trends is not confounded by biased results, we will collect duplicate samples and field measurements at the rate of approximately once every 10 samples. We will assess the relative percent difference between duplicate samples or measurements and flag data that do not meet the QA/QC guidelines detailed in SOP #12. Those data that do not meet the QA/QC stipulations may or may not be used in analyses, on a case by case basis. For example, parameters measured at low concentrations (e.g., less than five times the method detection limit) may be accepted, and both duplicates used in analysis. If duplicate samples meet the QA/QC guidelines, we will use the mean value of the duplicate measurements in data analyses. Explanations of how duplicates are handled in data analyses will be included in reports.

9.5.4 Other Analytical Considerations

For trend analyses performed using regression, trend will be investigated using a linear relationship. If analyses suggest a non-linear temporal pattern, serial autocorrelation, or lagged response to stressors, appropriate analytical modifications will occur. Adopting the philosophies of the precautionary principle (United Nations 1992) and the safe minimum standard of conservation (Ciriacy-Wantrup 1952; Berrens et al. 1998), the GLKN monitoring program seeks to identify potential natural resource problems early, before deleterious or irreversible ecosystem changes occur. Consequently, for most analyses that use the null hypothesis of no change through time, we will adopt $\alpha = 0.10$. Furthermore, for particularly variable parameters that are measured at relatively few stations, magnitudes of change that seem biologically meaningful (Johnson 1999) yet have $0.10 < \alpha < 0.20$ may nonetheless merit either increased attention (e.g., more extensive sampling) in future monitoring or experimental study. If water-quality monitoring tracks a relatively large suite of variables, Bonferroni or modified Bonferroni corrections (e.g., Hohm's method) may be performed, to maintain the 'familywise' alpha at 0.10. This is because, if 20 ecological variables were monitored at an alpha of 0.10, two of them would be expected to apparently exhibit 'significant' trends, when none of them had any actual trend. Roback and Askins (2005) argue, however, that if the main goal is simply initial screening for conservation problems or generation of hypotheses to be tested further (but not detection of real differences or trends), then alpha can be left at its standard comparison wise significance level, to avoid Type II errors.

Lotic systems are dynamic when compared to their more static lentic counterparts; water moves through the river network relatively quickly, with a finite residence time. Therefore, measurements

taken during the year at a particular station would constitute temporal subsamples rather than replicates (unless the domain of interest is only within that particular year, in which case the measurements would be independent, repeated measures). Repeated samples at a particular location would be likely to exhibit less correlation with each other than to samples at other points in the domain sampled at the same time. There are, however, analytical frameworks (e.g., repeated measures) that can accommodate multiple samples taken from the same location.

When the dataset contains non-detects, we will follow Helsel (2005) in conducting statistical analyses. Several methods exist for handling non-detects, each with advantages and drawbacks. Early in our program we will likely use a substitution method, where a value between zero and the detection limit will be substituted for the non-detect. The main drawback of this method is that estimates of the true variability are not possible. When the program is more mature and we have adequate data to determine the distribution of data, we will likely use maximum likelihood estimation, which works well for large sample sizes. Other methods, such as “regression on order statistics” and the non-parametric Kaplan-Meier method will be considered, and in consultation with a statistician, the most appropriate method for the data will be employed.

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Standard Operating Procedure #10: Reporting

Version 1.1

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Revision History Log

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Previous Version Number*	Revision Date	Author (with title and affiliation)	Location in Document and Concise Description of Revision	Reason for Change	New Version Number
1.0	2/11/2016	David VanderMeulen, Rick Damstra, GLKN	Updated the NPS contact author and authorship, distinguished between routine data summaries and periodic summary reports, added language about the use of web-based visualization and data summary tools, added resource briefs and newsletters as subsections under 10.4 Other Communications, fixed a handful of grammatical errors throughout	To clarify and update SOP based on changes that have occurred since Version 1.0 was published in 2008	1.1

*Add rows as needed for each change or set of changes tied to an updated version number

10.0 Introduction

A primary goal of the NPS Inventory and Monitoring Division is to ensure that the results and knowledge gleaned from monitoring are shared with all appropriate parties, especially the parks and their natural resource managers. Because the Network's main focus is to assist parks with monitoring needs, we will strive to provide park managers with clear, meaningful products to convey our findings.

While the Network primarily addresses concerns of the parks, its monitoring program has the potential to serve a much broader community. For example, monitoring projects can provide a starting point for external scientific research (especially to establish cause-effect relationships), and can provide insights for adaptive management on other public lands. The Network is also accountable to multiple organizations within the federal government, including the NPS I&M Division and the U.S. Congress. To ensure accountability and to meet the requests of all parties, we will provide the types of reports and communications detailed below.

10.1 Summary Reports

Routine data summaries will be completed annually for parks that are sampled within that year. The summaries will be compiled from data that has been uploaded to the EPA's STORET database and Aquatic Informatics Aquarius database. Hard copy or web-based summary reports will be produced periodically, with the primary audience being park resource managers.

Web-based summaries will primarily be in the form of visualization tools that allow the user to query data to create customized water quality summaries, which are currently under development. Hard copy summary reports will document our efforts and convey the findings of previous field seasons. At a minimum they will provide:

- a brief introduction that describes why that Vital Sign is being monitored;
- an outline of the sampling strategy, including the number of sites sampled, parameters measured, and analyses performed;
- data summaries, including tables and figures to enhance visual presentation, as well as a text explanation of the findings;
- any other relevant or significant findings; and
- a limited discussion section in which important results are interpreted.

The project manager (aquatic ecologist) will take the lead in writing the report, coordinate an internal review, and take responsibility for providing the report to parks.

10.2 Analysis and Synthesis Reports

Detailed reports in which data are analyzed and synthesized will be produced on a periodic basis, with the frequency depending on the given Vital Sign. The first analysis and synthesis reports will be written after at least three years of water quality data have been collected at a given park. The

frequency of subsequent detailed reports will depend on the data and whether or not trends seem to be occurring. As water resources at the parks are monitored repeatedly, in-depth analyses will be conducted for each park as well as across parks.

The reports will be written in the format of a scientific journal article (abstract, introduction, methods, results, discussion, literature cited) and will contain in-depth analyses as outlined in the protocol narrative and SOP #9, Data Analysis. Further, these comprehensive reports will:

- place the observed results in both a regional and historical context by relating them to other published literature;
- discuss the significance of the results in terms of environmental change; and
- provide management recommendations based on the findings.

The project manager will take the lead in writing the analysis and synthesis reports, and will coordinate an internal review. The target audience of these reports will be the parks (primarily the natural resource managers), the Network, and both regional and Servicewide I&M. Outside of the park service, the target audience includes the four state departments of natural resources (Indiana, Michigan, Minnesota, Wisconsin), the Minnesota Pollution Control Agency, the St. Croix River Interagency Basin Team, and the broader scientific community.

Drafts of analysis and synthesis reports will be reviewed internally and possibly sent to outside sources for further review, depending on how analytically complicated the methods are and on the gravity of the implications and recommendations.

10.3 Scientific Journal Articles

Because the large rivers protocol has been designed with rigorous standards for sampling design and analysis, monitoring results are expected to be highly defensible and meet the standards of the peer-review process. The publication of monitoring results in scientific journals will allow the Network to reach the scientific community in a way that internal NPS reports cannot. Further, peer-reviewed publications can promote collaborative investigation by members of the scientific community, either independently or in cooperation with the Network. Ultimately, this process should foster a greater understanding of ecosystem components and processes.

For these reasons, the Great Lakes I&M Network will strive to publish analysis and synthesis reports in peer-reviewed scientific journals. We will encourage the preparation of manuscripts by having reviewers of analysis and syntheses reports recommend whether publication is warranted and suggest appropriate journals. The aquatic ecologist and Network coordinator will track these recommendations and encourage and provide work time, respectively.

10.4 Other Communications

While reports are a definitive method of documenting the progress of each program, other means of communication can further disseminate information to a broader audience. To this end, we will provide the following additional types of communications:

10.4.1 Presentations to Park Natural Resource Managers

The project manager will periodically present the findings from the water quality monitoring program from previous years to the natural resource managers from the parks where monitoring took place. These presentations, which will likely occur at the annual technical committee meeting in March, will provide a concise synopsis of monitoring results as well as management considerations.

10.4.2 Conference Presentations

When possible, the project manager will present monitoring results at regional and national scientific conferences. Such presentations will allow the Network to reach the broader scientific community, as well as land managers and conservation practitioners. Potential conferences include those sponsored by the National Water Quality Monitoring Council, American Water Resources Association, the University of Minnesota Water Resources Center, the NPS Water Professionals Meeting, and the George Wright Society. At a more local scale, the Western Great Lakes Research Conference, which is sponsored in part by the Network, is a valuable venue for information exchange.

10.4.3 Resource Briefs

The project manager and assistant project managers will periodically publish short (typically one double-sided page) resource briefs (i.e., fact sheets) related to the GLKN aquatics program. These briefs, which are written for a lay audience, will convey key findings without using overly technical terminology or jargon. The briefs can be specific to a park (i.e., “Water Quality Monitoring at St. Croix National Scenic Riverway”) or topical (i.e., “Monitoring North Manitou Island”). Hard copies of resource briefs are disseminated to parks, park partners, and other collaborators, and provided electronically on GLKN’s website.

10.4.4 Newsletter Articles

GLKN publishes a newsletter titled *The Current* in the spring and fall, in which the project manager provides plans for water quality monitoring and field season summaries, respectively. The project manager will also occasionally write water quality related feature articles for the newsletter. Hard copies of the newsletter are disseminated to parks, park partners, and other collaborators, and provided electronically on GLKN’s website.

Standard Operating Procedure #11: Post-Season Procedures

Version 1.1

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Previous Version Number*	Revision Date	Author (with title and affiliation)	Location in Document and Concise Description of Revision	Reason for Change	New Version Number
1.0	2/11/2016	David VanderMeulen, Rick Damstra, GLKN	Updated the NPS contact author and authorship, added language about long-term storage of multiprobes, noted that equipment must be checked to ensure that it is clearly marked, added language about storing and inventorying equipment with GLKN and/or NPS property numbers, referenced chemical inventory records, replaced "Material Safety Data Sheets" with "safety data sheets", fixed a handful of grammatical errors throughout	To clarify and update SOP based on changes that have occurred since Version 1.0 was published in 2008	1.1

*Add rows as needed for each change or set of changes tied to an updated version number

11.0 Introduction

The proper maintenance and storage of field and laboratory equipment will prolong the life of the gear as well as simplify start-up procedures for the next field sampling season.

11.1 End of Season Procedures

11.1.1 Field Instrumentation and Equipment

When water quality probes are to be stored for extended periods of time, make the following preparations:

- Thoroughly clean the sensors.
- Remove installed batteries.
- Fill the storage cap approximately one-fourth full of tap water or pH 4.0 calibration solution. Store in a temperature-controlled environment. Do not expose sensors to freezing temperatures. Store away from direct sunlight. Although the instrument should be able to be reliably reactivated for field use with a minimum of effort before field use, it should be checked out well in advance of scheduled surveys to allow time for repair or replacement.

Refer to equipment manuals for more detailed instructions regarding maintenance of multiprobe sondes or other sensors.

End-of-season care of other equipment includes the following:

- Inspect and clean all equipment following the procedures detailed in SOP #5, Decontamination of Equipment to Remove Exotic Species. This effort minimizes the potential for transferring nuisance species from contaminated lakes to uncontaminated lakes.
- Lay out all ropes and sampling lines to dry completely, then coil or roll back into their holders.
- Store field equipment in protective storage cases to avoid damage.
- Clean all sample collection and storage containers in a 0.1N HCl acid bath followed by deionized water rinses as per SOPs #6 and #7.
- Ensure that all equipment is clearly marked (e.g., GLKN WQ).
- Return all of the equipment and supplies to the proper storage area. Keep them organized so they can be inventoried using the equipment and supply checklists, and for annual property inventories for equipment assigned a GLKN or NPS property number.
- Store calibration standards and electrolyte solutions in a temperature-controlled environment.
- Properly dispose of all chemical waste material.

11.1.2 Laboratory Equipment and Supplies

- Clean all lab ware in a 0.1N HCl acid bath followed by deionized water rinses as per SOP #7.

- Inventory all supplies, including chemical inventory records, and if necessary replace chemical and non-chemical supplies as soon as possible.
- Ensure that safety data sheets (formerly known as material safety data sheets) are up-to-date, complete, and readily accessible.

11.2 Data Management

There is no substitute for complete and accurate record keeping of field-derived data. Field technicians, crew leaders, and project leaders share responsibility for collecting, verifying, and documenting data according to the guidelines in this monitoring protocol and all applicable standard operating procedures. Refer to the GLKN Data and Information Management Plan (Hart and Gafvert 2006) for overall guidance and follow SOP #8, Data Entry and Management and SOP #10, Reporting, for additional details.

11.3 Quality Assurance/Quality Control

Taking proper care of all field and lab instrumentation and sampling gear is a fundamental part of any QA/QC program. Sensors that are properly cared for will likely be less variable, equilibrate more quickly in the field, and less prone to malfunction or failure. See SOP #12 for additional details on QA/QC procedures.

11.4 Literature Cited

Hart, M., and U. Gafvert (editors). 2006. Data management plan: Great Lakes Inventory and Monitoring Network. Great Lakes Inventory and Monitoring Network Report GLKN/2006/20. National Park Service, Ashland, Wisconsin. Available at http://science.nature.nps.gov/im/units/glkn/assets/docs/GLKN_SD8_Data_Management_Plan.pdf

Standard Operating Procedure #12: Quality Assurance/Quality Control

Version 1.2

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Revision History Log

The following table lists all edits and amendments to this document since the original publication date. Information entered in the log must be complete and concise. Users of this standard operating procedure will promptly notify the project manager and/or the Great Lakes Network (GLKN) data manager about recommended and required changes. The project manager must review and incorporate all changes, complete the revision history log, and change the date and version number on the title page and in the header of the document file. For complete instructions, please refer to Revising the Protocol, SOP #13.

Previous Version #	Revision Date	Author (with title and affiliation)	Location in Document and Concise Description of Revision	Reason for Change	New Version #
1.0	2/25/2010	David VanderMeulen, Joan Elias, Rick Damstra, GLKN	Table 4 – update detection levels required; info on optical DO probes added; Table 9 – clarify & update calibration frequency	To bring SOP into alignment with procedures practiced	1.1
1.1	3/18/16	David VanderMeulen, Rick Damstra, GLKN	Updated the NPS contact author and authorship, added language about the use of web-based visualization and data summary tools, continuous monitoring with thermal arrays, data certification, and Aquarius database, updated monitoring sites in Table 1, added Attachment C, Sample Data Certification Form, fixed a handful of grammatical errors throughout.	To clarify and update SOP based on changes that have occurred since Version 1.1 was published in 2010	1.2

*Add rows as needed for each change or set of changes tied to an updated version number

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This document was prepared largely through the use of available National Park Service guidance and documents prepared for the Great Lakes Network Inventory and Monitoring program. The authors wish to acknowledge these sources and thank them for the valuable contribution that they have made to this document. Some verbiage contained within this standard operating procedure (SOP) was first written in the protocol narrative and other SOPs, and we wish to thank the authors of those sections.

12.0 Purpose

This standard operating procedure (SOP) defines procedures for quality assurance and quality control to be used with the Great Lakes Network protocol for monitoring water quality in large rivers.

Quality assurance is the planned and systematic pattern of all actions necessary to provide adequate confidence, or assurance, that a project outcome optimally fulfills expectations. Quality control is the systematic evaluation of the various aspects of a project to ensure, or control, that the standards of quality are being met. Quality control involves specific tasks undertaken to determine the reliability of field and laboratory data. Together, quality assurance/quality control (QA/QC) is a substantial part of any monitoring program. The objective of QA/QC is to ensure that the data generated by a project are meaningful, representative, complete, precise, comparable, scientifically defensible, and reasonably free from bias (Irwin 2006).

12.1 Applicability

Procedures in this SOP will be implemented during all work pertaining to monitoring water quality in large rivers as described in the protocol narrative. This SOP is designed to assure that all data obtained will contribute quality information to an understanding of the ecological integrity of park units of the Great Lakes Network.

The rationale for dividing a sampling protocol into a protocol narrative with supporting SOPs is based on the following:

- The protocol narrative is a general overview of the protocol that gives the history and justification for doing the work and an overview of the sampling methods, but does not provide all methodological details. The protocol narrative will only be revised if major changes are made, such as changes in sampling design.
- The SOPs are specific step-by-step instructions for performing a given task. They are expected to be revised more frequently than the protocol narrative.
- Usually, when a SOP is revised, it is not necessary to revise the protocol narrative to reflect the specific changes made to the SOP. All versions of the protocol narrative and SOPs will be archived.

The steps for changing the protocol (either the protocol narrative or the SOPs) are outlined in Procedures for Revising the Protocol (SOP #13).

12.2 Summary

Large rivers are important and valuable resources at two of the parks of the Great Lakes Network. Both the St. Croix National Scenic Riverway (SACN) and Mississippi National River and Recreation Area (MISS) were established to highlight and protect large river resources. Monitoring at MISS was suspended in 2014 after it was determined that water quality monitoring by MCES and other agencies was adequately characterizing water quality conditions throughout MISS (Lafrancois et al. 2013, VanderMeulen 2013). As a consequence, work at SACN was expanded to annual monitoring,

beginning in 2014 and continuing to present. The St. Croix and Namekagon rivers at SACN are used extensively by visitors for fishing, boating, swimming, and other recreational activities. The preservation of large river water quality and quantity is of utmost importance to park managers, researchers, and the general public. Monitoring basic water quality ranked among the highest of the Network's vital signs (Route 2004).

A national review panel assembled by the NPS Water Resources Division (WRD) recommended a suite of five parameters be measured across all NPS monitoring networks (NPS 2002). In addition to these five mandated parameters (temperature, pH, specific conductance, dissolved oxygen, and flow/water level) we added a measure of clarity (secchi depth or transparency depth) to our core suite. The core suite was ranked highest among potential vital signs for aquatic systems of GLKN parks, although it was recognized that these measurements were less diagnostic of water quality degradation than biotic communities and other water quality variables, such as turbidity, nutrients and chlorophyll-*a*, and ions.

Inputs of excess nutrients, invasion and spread of exotic species, and contaminants from atmospheric fallout and surface runoff, and how these stressors affect the chemical and biological functions of rivers are key issues of concern. By monitoring an advanced suite of parameters (turbidity, nitrogen and phosphorus species, dissolved organic carbon, major ions, alkalinity, dissolved silica, and chlorophyll-*a*), we will provide data for a thorough understanding of changes over time.

Our overall goal is to develop a program for monitoring water quality in large rivers that will contribute to an understanding of the health of ecosystems in park units of the Great Lakes Network. The monitoring protocol is intended to document water quality status and trends for individual river sites and provide an indication of status and trends on a park-wide basis. The protocol includes analysis of historical data, sample design, field and laboratory methods, data analysis and reporting, and training and operational requirements.

12.3 Definitions and Abbreviations

Term	Definition
Equipment blank	<i>Also known as field blank.</i> A sample of distilled, deionized water taken to the field, opened, and used as sample water would be (i.e., poured through equipment used to handle samples).
Trip blank	Used to indicate potential contamination due to migration of volatile organic chemicals (VOCs) from the air on the site or shipment into the sample vial. Consists of laboratory distilled, deionized water in a 40 mL glass vial sealed with a Teflon septum and is unopened in the field.
Laboratory calibration blank	Distilled, deionized water injected directly into an instrument; indicates contamination in instrument or source of water.
Laboratory reagent blank	<i>Also known as method blank.</i> Distilled, deionized water manipulated as if it were a sample (digestions, extractions, etc.).

Term	Definition
CCV	Continuing Calibration Verification
EPA	Environmental Protection Agency
EMAP	Environmental Monitoring and Assessment Program
GLKN	Great Lakes Inventory and Monitoring Network
m	Meter
MDL	Method Detection Limit
MISS	Mississippi National River and Recreation area (Minnesota)
mg/L	Milligrams per liter
ML	Method Limit of quantitation
MQO	Measurement Quality Objective
NAWQA	National Water Quality Assessment Program
NPS	National Park Service
NPSTORET	National Park Service Storage and Retrieval Database
QA/QC	Quality Assurance/Quality Control
QAPP	Quality Assurance Project Plan
Reference standard	Independently created solution of known concentration.
RPD	Relative Percent Difference
SACN	St. Croix National Scenic Riverway (Minnesota/Wisconsin)
SDS	Safety Data Sheet
SOP	Standard Operating Procedure
STORET	Storage and Retrieval Database
USGS	United States Geological Survey
µg/L	Micrograms per liter
WRD	Water Resources Division of the National Park Service

12.4 Personnel Responsibilities

Personnel will study this SOP prior to beginning work on the project and follow its procedures in order to conduct the project according to outlined QA/QC procedures. This will ensure consistency and comparability when changes in personnel occur.

The role of the **project manager** is to serve as a liaison among other related water quality monitoring projects conducted by partners (e.g., state monitoring programs), park staff, other Network staff (field personnel, data manager), contracted analytical laboratories, and other GLKN monitoring project managers. The individual will coordinate with resource management staff at the parks to ensure parks are informed of monitoring activities. Specific responsibilities of the project manager include the following:

- Coordinate field schedules and availability of supplies with field personnel
- Develop a training program for field personnel
- Develop, document, and oversee the implementation of standard procedures for field data collection and data handling
- Coordinate logistics with park staff
- Develop quality assurance and quality control measures for the project, supervise staff training and conduct quality assurance checks of field sampling techniques at least once, mid-season, with each field crew
- Contract with an analytical laboratory for analysis of water samples, ensure lab results meet program needs (e.g., QA/QC procedures, meaningful minimum detection limits for low level strength waters, adequate reproducibility of replicate samples)
- Supervise or perform data entry, verification, and validation
- Summarize and analyze data, prepare reports
- Serve as the main point of contact concerning data content

The project manager will also work closely with the data manager in the following capacities:

- Complete project documentation in NPSTORET and Aquarius (describing who, what, where, when, why and how of a project)
- Develop data verification, validation, and certification measures for quality assurance
- Ensure staff are trained in the use of database software and quality assurance procedures
- Coordinate changes to the field data forms and the user interface for project databases
- Identify sensitive information that requires special consideration prior to distribution
- Manage the archival process to ensure regular archival of project documentation, original field data, databases, reports and summaries, and other products from the project

- Define how project data will be transformed from raw data into meaningful information and create or oversee data summary procedures to automate and standardize this process
- Establish meaningful liaisons with state counterparts to promote sharing of data on a timely basis

The **field crew leader** will be responsible for preparing supplies and equipment for field season; ordering needed supplies; making travel arrangements; maintaining sampling equipment; maintaining field vehicles; ensuring field personnel follow sampling protocols; preparing field data forms, chain of custody forms, and analytical service request forms for each site; notifying contract laboratory of planned sample shipment date; and providing project manager with field trip and check-in schedule.

Field personnel should be adequately experienced and/or trained in using field and water quality sampling equipment. This experience is best obtained through a combination of classroom and hands-on training, including pilot-testing equipment at a nearby water body. Personnel should be familiar with the instruction manuals for all equipment, particularly with regard to calibration and maintenance procedures. See SOP #2, Training and Safety, for more details on training.

Field personnel will be responsible for following protocols and standard operating procedures during field activities and the handling of samples.

Roles of staff are explained in more depth in Section 5 of the large rivers protocol narrative, along with training and requirements.

12.5 Sampling Process and Design

Two major ecological monitoring programs (the USGS National Water-Quality Assessment (NAWQA) program and the EPA Environmental Monitoring and Assessment Program-Aquatic Resource Monitoring (EMAP-ARM)) use different site-selection methods. The GLKN goal is to combine the strengths of both programs for monitoring water quality of lotic systems in the NPS Great Lakes Network. For further information see the large rivers protocol narrative.

The information goals and statistical requirements determine the sampling frequency. To identify the long-term trends in a stream or river, the sampling frequency should be sufficient to identify a statistical trend beyond the background variability of a dynamic flow system. Therefore, it is essential to evaluate existing data prior to establishing 1) the number of sites and 2) the frequency of sampling. Statistical power analysis based on the power to detect change guides the selection of these two parameters, using calculations of statistical sample size and analysis of sensitivity to sampling frequency, respectively. The goal is to be able to detect 20% change at 80% power and 5% significance. Due to natural variability of the different parameters, the number of sampling sites and sampling duration time required to detect this level of change for all parameters of interest may not be possible within budget constraints.

12.5.1 Site Selection

This program includes both randomly-selected and nonrandomly-selected sites. All sites will be analyzed on an individual basis, to determine water quality trends at each location. Only the randomly-selected sites will be grouped for analysis of population statistics and trends.

GLKN monitoring site locations within MISS and SACN are shown in the protocol narrative. All five sites at MISS were nonrandomly-selected. At SACN, Six sites were randomly-selected, and since implementation of monitoring the number (and in some cases the location) of nonrandomly-selected sites has decreased from seven in 2007 to four in 2013 (to present) (Table 12-1). In summary, All site locations described in the protocol narrative were field-verified for representativeness and accessibility before inclusion in the monitoring design.

Table 12-1. Number of randomly and nonrandomly-selected monitoring site at MISS and SACN in 2016.

Station Type	MISS	Upper SACN	Lower SACN
Random	--	3	3
Nonrandom	5	4	--
Total	5	7	3

12.5.2 Field Collection Parameters

Core field parameters will be measured with multiprobes or individual sensors. Samples will be collected monthly for nutrients, chlorophyll-*a*, and total suspended solids (a measure of turbidity) during the open water season, which is usually April/March through November, monitoring spring, summer, and autumn flows. The core water quality suite (dissolved oxygen, temperature, specific conductance, pH, flow, and clarity) will also be monitored monthly. Water samples for laboratory analysis of major cations and anions, alkalinity, dissolved silica, and dissolved organic carbon will be collected quarterly, or three times during the open water season. In addition, a near-bottom sample for TP will be collected once annually from thermally stratified sites. Randomly-selected and nonrandomly-selected stations in each park unit will be sampled on the same frequency and at the same time each sampling period.

Collect water sample(s) with an integrated sampling tube or Van Dorn and process as per SOP #7. In the field log book and on the field data sheet, record information related to the sample collection, including:

1. Site name and site identification code
2. Sample date, time, and depth.
3. The amount of water collected.
4. Whether duplicate samples for quality control were collected at this site.
5. Any additional notes or observations pertinent to this sample or location for this sampling period.

Additionally, always keep in mind the following:

- Sample containers should be labeled in indelible ink with, at a minimum, the station name, date and time of collection, and preservation method, if applicable.
- Follow all SOPs for sample collection and preservation.
- To ensure the integrity of the sample, be aware of possible sources of contamination. Contamination introduced during each phase of sample collection and processing is additive and usually is substantially greater than contamination introduced elsewhere in the sample handling and analysis process.
- Use appropriate procedures and quality-assurance measures that ensure sample representativeness and integrity and that meet study criteria. The degree to which a sample can be considered representative of a water body depends on many interrelated factors including temporal and spatial homogeneity of the water body, sample size, and the method and manner of sample collection.

12.5.3 Field Analysis

For discrete monitoring dissolved oxygen, temperature, specific conductance, and pH will be measured in the field using a multiprobe and following the methods listed in Table 12-2. Multiprobes typically perform within the specifications detailed in Table 12-3. Other parameters may be analyzed at the field station by field personnel in the future.

Specification of quantification ability for field multiprobes and sensors is not a straightforward data quality objective exercise. Most field parameters tend to be those that characterize the waterbody and are not usually based on a criterion limit as would be used for a toxic pollutant. Therefore, instrument selection should be based on the parameters and ranges they can measure, but instrument-specific estimations of the range of uncertainty for each parameter will have to be made when interpreting data. Details on estimating instrument sensitivity are included below in section 12.8.2.

Table 12-2. Core suite of field variables and required in situ measurement method.

Field Variable	Method
Temperature (°C)	EPA 170.1
pH	EPA 150.1
Specific conductivity (µS/cm)	EPA 120.1
Dissolved oxygen (mg/L)	EPA 360.1
Clarity (cm)	Transparency tube

Table 12-3. Typical sensor performance specifications for multiprobe field instruments.

Sensor	Expected Range	Reporting Resolution*	Estimated Bias	Stabilization Criteria
Temperature	-5°C to 45°C	0.01°C	±0.15°C	Thermistor: ±0.2°C Glass: ±0.5°C
Specific Conductivity (SC25)	0 to 2000 µS/cm	µS/cm (range dependent)	±0.5% of reading +1 µS/cm	≤100 µS/cm: ±5% >100 µS/cm: ±3%
pH	1 to 14 units	0.01 unit	±0.2 units	±0.1 standard unit
Dissolved Oxygen (Conc.)	0 to 50 mg/L	0.01 mg/L	0 to 20 mg/L: ±0.2 mg/L 20 to 50 mg/L: ±0.6 mg/L	±0.3 mg/L
Dissolved Oxygen (% sat.)	0%–200%	0.1%	ca. ±2%	±2%
Depth – Z (pressure sensor)	0–>100 m	0.1 m	ca. 0.1 m	0.1 m

* Resolution specifications are supplied by the manufacturers of the measuring meters. They are not necessarily closely related to real-world (outdoor) precision or bias, and are sometimes more related to the number of significant figures reported rather than how accurate the extra significant figures are. This is why we will control measurement sensitivity in the actual outdoor measuring environment at least once a year by calculating alternative measurement sensitivity (AMS; see Irwin 2006 for more details on AMS)

12.5.4 Laboratory Analysis

Samples will be collected in the field according to SOP #6 for shipment to a contract laboratory that meets QA/QC requirements outlined in Section 12.8.4. Samples will be analyzed for nutrients (TP, TN, NO₃+NO₂-N, NH₄-N), total chlorophyll-*a*, and total suspended solids monthly, while alkalinity, major ions (Cl⁻, SO₄²⁻, Ca²⁺, Na⁺, K⁺, and Mg²⁺), DOC, and SiO₂, which tend to be less variable, will be monitored quarterly.

Methods chosen will meet quantification limits according to the criteria shown in Table 12-4. The method limit of quantitation (ML) of the chosen method should be two to ten times lower than the

typical expected low value. The ML is the lowest value that can be quantified with certainty. The value in the “ML needed” column of Table 12-4 is two to ten times lower than the lowest value found in the pilot year of sampling water quality for each analyte. This estimation is used as guide to the selection of the method that will be needed for this project.

Table 12-4. Analytical detection levels required for GLKN water quality monitoring.

Analyte	Typical Low Value Found	Criteria Source ^{a, b}	Comment	ML Needed
Alkalinity (mg/L) ^c	ND (<1)	Lowest value pilot year	MDL unknown	--
DOC (mg/L) ^c	3	Lowest value pilot year	Lab MDL 0.1	0.3 mg/L
Ca ²⁺ (mg/L) ^c	2.4	Horizon report	MDL unknown	0.24 mg/L
Cl ⁻ (mg/L) ^c	0.273	Lowest value pilot year	Lab MDL 0.025	0.02 mg/L
Chl-a (µg/L)	0.63	EPA	Lower of VII and VIII	0.06 µg/L
Mg ²⁺ (mg/L) ^c	0.47	Horizon report	MDL unknown	0.04 mg/L
K ⁺ (mg/L) ^c	0.24	Lowest value pilot year	Lab MDL 0.06	0.02 mg/L
Na ⁺ (mg/L) ^c	0.5	Horizon report	MDL unknown	0.05 mg/L
SO ₄ ²⁻ (mg/L) ^c	0.625	Lowest value pilot year	Lab MDL 0.025	0.06 mg/L
TP (µg/L)	10	EPA	Lower of VII and VIII	1.0 µg/L
TN (µg/L)	0.38	EPA	Lower of VII and VIII	0.03 mg/L
NH ₄ -N (µg/L) ^c	5.0	Horizon report	MDL unknown	1.0 µg/L
NO ₃ +NO ₂ -N (µg/L) ^c	2.0	Horizon report	MDL unknown	1.0 µg/L
SiO ₂ (mg/L)	ND (<0.1)	Lowest value pilot year	Lab MDL 0.2	0.1 mg/L

NOTES:

^a: EPA = EPA Ecoregional nutrient criteria recommendations, lower value recommended for aggregate ecoregions VII and VIII (USEPA 2000).

^b: WQS = criteria value listed in Ledder (2003) state water quality standards

^c: parameter has no official criterion for any of the states, the value reported in the analytical requirement column is the lowest value determined in the pilot year for rivers or lakes or the lowest value found in a spot check of Horizon reports for each park.

The majority of the parameters listed in Table 12-4 are used in waterbody characterization and do not have criterion lower limits as would a toxic pollutant. Some of the criteria available are listed as maximums not to be exceeded and, as such, do not guide the selection of a method by ML.

Guidelines to be used, therefore, include relevant state water quality standards, U.S. Environmental Protection Agency (EPA) ecoregional nutrient recommendations (USEPA 2000), state nutrient criteria as they are developed, data results for these parameters from the pilot studies carried out by GLKN, lowest values for the parameter found in relevant Horizon reports (NPS 1995a, 1995b, 1995c, and 1999), and/or values based on the best professional judgment of GLKN staff.

12.6 Sampling Methods

Consistent methods are important to long-term quality data. In actuality, the ideal conditions are not always met in the field or in the lab and changes in staff do occur. Therefore, documentation of procedures, site conditions, laboratory analysis, and reasons for deviations of any kind is important in and of itself for long-term projects. Personnel will be encouraged to write down more than they feel may be necessary in the moment as the future interpretation of their data will depend on the written record and not the memory of an individual.

12.6.1 Site Locations

Refer to description of monitoring station location, directions, and photos to verify correct location. Verify coordinates with a GPS unit. Document this verification. This information will be contained in a field binder along with a table of previous field measurements to compare with new measurements, and a discharge rating curve to verify new stage readings.

Whenever possible, collect water samples from the centroid of flow, in a vertical profile of the water column. The centroid of flow is the midpoint of streamflow, or the point on the river's cross-section perpendicular to flow that has equal volumes of flow to the right and to the left sides. Sampling from the centroid of flow assures that average water quality conditions are being sampled, and that water quality results are reasonably representative of average flow conditions at the particular site being sampled.

Collect samples on the upstream side of the boat, to minimize contamination and disturbance. Avoid surfactants, floating debris, and turbid aeration when collecting samples. Rinse the integrating sampler and compositing jug three times with surface water before taking samples. Discard rinse water or excess sampling water on the downstream side of the boat. Allow any disturbed sediments to clear before taking the actual sample. Always follow procedures for minimizing contamination during the sampling process (See SOPs #5 and 6 for more details).

12.6.2 Field Water Measurements

Before making field measurements during discrete monitoring, properly-calibrated sensors (see SOP #6) must be allowed to equilibrate to the condition of the water being monitored. Sensors have equilibrated adequately when instrument readings have stabilized, that is, when the variability among measurements does not exceed an established criterion. The criteria for stabilized field readings were defined operationally by O'Ney (2005) for a set of three or more sequential measurements (Table 12-5). The natural variability inherent in surface water or ground water at the time of sampling generally can be compared with these stability criteria and indirectly relates to the short term bias or the long term accuracy that should be attainable with a calibrated instrument. Dissolved oxygen sensors that

utilize a Clark cell membrane may require up to five minutes to stabilize, while optical dissolved oxygen sensors and sensors for the other parameters typically require less than 60 seconds to stabilize. Observers should only note instrument readings after the stabilization criteria in Table 12-5 are met for all the sensors, and then record readings for all parameters at once.

Measure field water quality variables and conduct sampling according to SOP #6, field measurements and water sample collection. Quality results require consistent measurement methods and detection limits.

Depth profiles of temperature, specific conductance, pH, and dissolved oxygen will be measured at each sampling station using a multiparameter instrument (multiprobe). Clarity will be measured using a transparency tube and Secchi disk (on sites where possible). These core parameters will be measured when water samples for analysis of the advanced parameters are collected.

Begin just below the water's surface (ca. 0.3 m depth) and take readings after stabilization of the multiprobe. Lower the sensors to collect a vertical profile of field parameters at 1m intervals until 20 m depth, and then every 2–5 m depending on overall depth and gradients. Replicate 10% of the readings (e.g., at 1 m, 10 m, etc.); take the replicate readings immediately following the original readings. Values should agree within 10% or the acceptance criteria in Table 12-5, whichever is larger.

Table 12-5. Recommended instrument stabilization criteria for recording field measurements^{a,b}.

Standard Direct Field Measurement	Stabilization Criteria (O'Ney 2005)	Stabilization Criteria <i>In situ</i> Multisensors (WOW 2005)
Temperature^c:		
Thermistor Thermometer	± 0.2°C	± 0.2°C (5%)
Liquid-in-glass Thermometer	± 0.5°C	
Specific Conductivity (SC25)^d		
When ≤ 100 µS/cm	± 5%	<5 µS/cm (10%)
When > 100 µS/cm	± 3%	
pH^e: Meter displays to 0.01	± 0.1 unit	± 0.2 unit (10%)
Dissolved oxygen^e:		
Amperometric (same as polarigraphic) method	± 0.3 mg/L	± 0.5 mg/L (10%)

^a: Resolution/sensitivity is a data quality indicator related to detection limits but typically handled differently for field probes than for laboratory parameters. For more information, see Irwin (2004).

^b: In the case of field probes, bias is typically a best case maximum deviation from known correct values (typically based on comparisons with known NIST certified reference materials or standards). True accuracy is a combination of high precision and low bias, and is hard to quantify with the small sample sizes used to control bias and precision in typical field measurements (see Irwin 2004 for more details).

^c: Recommended sensor calibration is quarterly.

^d: Recommended sensor calibration is daily.

^e: Sensor calibration at beginning of sampling day with a calibration check at the end of day.

At all sites, record visual observation information required on the data sheet. Such observational data can provide important information to the interpretation of field measurements.

If any analyses are to be done in-house, conduct these analyses as soon as possible upon return from the field. A clean analytical station should be prepared in which to work, free of food items, mud, lubricants, or lab chemicals. Hands should be thoroughly washed.

12.6.3 Samples for Laboratory Analysis

Prepare bottles and labels prior to field sampling as per SOP# 6. Collect samples with an elbow-deep grab or using a 0–2 m integrating tube sampler as per SOP #6. For all sampling, it is critical to avoid sampling water showing evidence of oil, gasoline or anything else from the boat. It is best to kill the engine and set the anchor, if possible, although this may not be possible or advisable in bad weather or with a balky engine.

Prior to filling sample bottles sent by the laboratory, first rinse the bottle once with sample water, if the bottle is not pre-preserved. Some analyses require preservation of the sample with acid, which may be added to the bottle by the contract laboratory; in these cases do not rinse the bottle first. Take care not to overfill the bottle if the bottle is pre-acidified, as overfilling will flood the acid out of the bottle. If the samples requiring acid preservation are not pre-preserved, use caution to add the ampule of preservative supplied by the lab and immediately rinse hands in water if acid is spilled. Protective gloves are recommended.

Table 12-6 summarizes the variety of methods, detection limits, preservation techniques, and holding times for water samples addressed by this protocol. These methods conform to those used by Minnesota, Wisconsin, and Michigan for state certification of environmental laboratories involved in Clean Water Act or drinking water sample analysis (MDH 2005, WSLH 2003, MDEQ 2005). They are also used by EPA-funded research projects of natural waters in the upper Midwestern U.S. Holding times shall in no case be less stringent than those recommended by EPA in 40 CFR Part 136 to 136.3 and appendices. Refer to SOP #6 for additional details regarding sample collection and preservation.

Samples are stored in a cooler with ice packs during field sampling. Prepare samples for shipment according to the contract laboratory's protocols. These protocols will be provided to the field sampling personnel for each sampling round so that the proper procedures are accessible in the event of contract laboratory changes. In general, samples are shipped on ice to maintain a temperature of approximately 4°C. A plastic bag is placed in the cooler first. Sample bottles are packed among zip lock bags of ice and/or ice packs to prevent water leakage into the sample bottle during shipment. Prepare a temperature check bottle for each anticipated cooler, if recommended by the contract analytical laboratory. Use tap water to fill an extra bottle of the same size used for one of the analytes and label as "Temperature Check." Store this check bottle in refrigerator with other samples; package and send to the analytical laboratory with the other samples. Some analyses require the sample to be frozen for shipment; such frozen samples are likewise packed among ice packs for shipment to the contract laboratory.

Table 12-6. Example range of analytical methods, method detection limits (MDLs), containers, and holding times.

Analyte	Analytical <i>Note 1</i>	Method #	Det. Limit	Vol. (ml)	Filter	Preservation	Sample Bottle <i>Note 2</i>	Hold Time
Alkalinity	Titrimetry	310.1 EPA-NERL	10 mg/L	--	--	4°C	--	14 d
	Spec. auto.	310.2 EPA-NERL	10 mg/L	--	--	4°C	--	14 d
	Titrimetry	NFM USGS-OWQ	0.01 meg/L	--	<i>Note 4</i>	None	--	None
Ca ⁺²	ICP	3120B APHA	10 ug/L	--	<i>Note 3</i>	pH<2 HNO ₃	P or G	6 mo
	Titrimetry	215.2 EPA-NERL	0.5 mg/L	--	<i>Note 3</i>	4°C	--	6 mo
	FAA	I-3152 USGS-NWQL	0.1 mg/L	250 mL	<i>Note 3</i>	pH<2 HNO ₃	P	180 d
Cl ⁻	IC	300.0 EPA-NERL	0.02 mg/L	--	--	4°C	P or G	28 d
	Colorimetry	325.2 EPA-NERL	1 mg/L	--	--	4°C	--	28 d
	Titrimetry	4500-Cl APHA	0.15 mg/L	100 mL	--	4°C	P or G	28 d
Chlorophyll-a	Spect.	10200 APHA	2 ug/L	≤1 L	<i>Note 4</i>	Freeze filter	P	30 d
Color	Spect.	110.2 EPA-NERL	5 Pt units		<i>Note 5</i>	4°C	G	48 hr
	Vis. Comp.	I-1250 USGS-NWQL	1 Pt-co	250 mL	<i>Note 5</i>	4°C	P	30 d

Source: National Environmental Methods Inventory website (NEMI 2006)

This list is not an endorsement of any particular method or laboratory for any particular analyte. Rather it is to be used as a reference for the range of analytical methods available for each analyte. There are surface water conditions (pH, turbidity, other elements) that make a particular method unsuitable for a particular situation. As GLKN is monitoring surface water, the methods listed were chosen as representative of the lower range of detection limits.

Note 1. CIE-UV= capillary ion electrophoresis with UV detection, FAA = flame atomic absorption, FIA = flow injection analysis, IC= ion chromatography, ICP = inductively coupled plasma, Spec. auto = spectroscopy with autoanalyzer

Note 2. P = plastic (polypropylene), G=glass, AG=amber glass

Note 3. 0.45µm membrane filter. Pre-filter for dissolved portion analysis.

Note 4. 0.45µm glass fiber filter.

Note 5. 0.45µm membrane filter or centrifuge is recommended to remove suspended solids that affect color, however some color will also be removed.

Note 6. Patton, C. J., and J. R. Kryskalla. 2003. Evaluation of alkaline persulfate digestion as an alternative to Kjeldahl digestion for determination of total and dissolved nitrogen and phosphorus in water. U.S. Geological Survey Water-Resources Investigations Report 03-4174. <http://nwql.usgs.gov/Public/pubs/WRIR03-4174/WRIR03-4174.html>

Table 12-6 (continued). Example range of analytical methods, method detection limits (MDLs), containers, and holding times.

Analyte	Analytical Note 1	Method #	Det. Limit	Vol. (ml)	Filter	Preservation	Sample Bottle Note 2	Hold Time
DOC	Spect.	415.3 EPA	0.018 mg/L	125	Note 3	pH<4 H ₂ SO ₄	G	28 d
	Spect.	0-1122-92 USGS	0.1 mg/L	--		4°C	AG	
K ⁺	ICP	3120B APHA	0.3 mg/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
	FAA	3111B APHA	5 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
Mg ⁺²	ICP	3120B APHA	20 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
	FAA	3111B APHA	0.5 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
Na ⁺	ICP	3120B APHA	30 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
	FAA	3111B APHA	2 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
NH ₄ -N	Selective elec.	4500-NH ₃ E	0.08 mg/L	--	--	4°C/pH2, 0°C	--	24hr / 28d
	Colorimetry	350.2 EPA-NERL	0.08 mg/L	--	--	pH<4 H ₂ SO ₄	--	28 d
	Titrimetry	4500-NH ₃ APHA	5 mg/L	--	--	4°C/pH2, 0°C	--	24 hr / 28 d

Source: National Environmental Methods Inventory website (NEMI 2006)

This list is not an endorsement of any particular method or laboratory for any particular analyte. Rather it is to be used as a reference for the range of analytical methods available for each analyte. There are surface water conditions (pH, turbidity, other elements) that make a particular method unsuitable for a particular situation. As GLKN is monitoring surface water, the methods listed were chosen as representative of the lower range of detection limits.

Note 1. CIE-UV= capillary ion electrophoresis with UV detection, FAA = flame atomic absorption, FIA = flow injection analysis, IC= ion chromatography, ICP = inductively coupled plasma, Spec. auto = spectroscopy with autoanalyzer

Note 2. P = plastic (polypropylene), G=glass, AG=amber glass

Note 3. 0.45µm membrane filter. Pre-filter for dissolved portion analysis.

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Note 6. Patton, C. J., and J. R. Kryskalla. 2003. Evaluation of alkaline persulfate digestion as an alternative to Kjeldahl digestion for determination of total and dissolved nitrogen and phosphorus in water. U.S. Geological Survey Water-Resources Investigations Report 03-4174. <http://nwql.usgs.gov/Public/pubs/WRIR03-4174/WRIR03-4174.html>

Table 12-6 (continued). Example range of analytical methods, method detection limits (MDLs), containers, and holding times.

Analyte	Analytical Note 1	Method #	Det. Limit	Vol. (ml)	Filter	Preservation	Sample Bottle Note 2	Hold Time
SiO ₂	ICP	3120B APHA	20 ug/L	--	Note 3	pH<2 HNO ₃	P or G	6 mo
	Spect.	4500- SiO ₂ D APHA	0.04 mg/L	--	Note 3	No, 4°C	P	28 d
	FIA-Spect.	4500- SiO ₂ F APHA	0.78 ug/L	--	Note 3	No, 4°C	P	28 d
TSS	Gravimetric	I-3765 USGS-NWQL	1 mg/L	250–500	Note 4	4°C filter	P	NA
SO ₄ ⁻²	IC	4110C APHA	75 ug/L	--	Note 3	pH<4 H ₂ SO ₄	P or G	--
	CIE-UV	D6508 ASTM	0.1 mg/L	--	Note 3	pH<4 H ₂ SO ₄	--	ASAP
	Spect.	37512 EPA-NERL	0.5 mg/L	--	Note 3	pH<4 H ₂ SO ₄	P or G	28 d
TP	Spect.	I-2606 USGS-NWQL	0.001 mg/L	125 mL		MgCl 4°C	BrownP	30 d
	Alkaline P	USGS 2003	0.01 mg/L	120 ml	Note 6	4°C /H ₂ SO ₄	--	48 hr / 30 d
	ICP	200.7 EPA-NERL	60 ug/L			pH<2 HNO ₃	P	6 mo
TN	Alkaline P	USGS 2003	0.03 mg/L	120 ml	Note 6	4°C /H ₂ SO ₄	--	48 hr / 30 d
	Titrimetry	4500-N	0–100 mg/L	--	--	4°C	AG	7 d
	Combustion	440.0 EPA-NERL	0.1 mg/L	--	--	Filter	--	100 d

Source: National Environmental Methods Inventory website (NEMI 2006)

This list is not an endorsement of any particular method or laboratory for any particular analyte. Rather it is to be used as a reference for the range of analytical methods available for each analyte. There are surface water conditions (pH, turbidity, other elements) that make a particular method unsuitable for a particular situation. As GLKN is monitoring surface water, the methods listed were chosen as representative of the lower range of detection limits.

Note 1. CIE-UV= capillary ion electrophoresis with UV detection, FAA = flame atomic absorption, FIA = flow injection analysis, IC= ion chromatography, ICP = inductively coupled plasma, Spec. auto = spectroscopy with autoanalyzer

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Follow the shipping company's requirements in preparation of the cooler of samples for shipment. Packaging problems may cause delays in shipment, which can mean that samples do not arrive at the laboratory at the proper temperature or past holding time, compromising data quality. In general, the ice and samples should be contained within a sealed plastic bag within the cooler so that the cooler does not leak. Packing tape should be wrapped around the cooler vertically and around the lid seam horizontally to ensure it remains closed. A note on the cooler that it contains water samples is helpful as well so that if leakage occurs, carriers will know the contents are not hazardous.

12.6.4 Sampling Forms

Before leaving the monitoring site, all field forms and sample labels must be reviewed for legibility, accuracy, and completeness. Any changes in procedure due to field condition must be explained in the comments section. Make sure the information is complete on all forms. Record the departure time on the field form. After reviewing each form, initial the upper right corner of each page of the form. Document any photos taken by including the photo number and roll number or digital camera photo number on the field form.

12.7 Handling and Custody

One part of proper data and sample handling procedures is to provide a complete record of the methods and procedures followed. Complete records are important to long-term monitoring so that anyone using the data may trace the sampling history.

12.7.1 Field Data

Field data will be collected on forms printed on waterproof paper. While at each monitoring site, the information recorded on the forms should include:

- Date and day of week
- Time of arrival
- Names of field team members
- GPS coordinates, to verify location
- Current weather (air temperature and wind speed) and relevant notes about recent weather (storms or drought)
- Observations of water quality conditions
- Description of any photographs taken
- Multiparameter sonde (model), calibration date, and field measurements of core suite variables
- Sample identification numbers and collection times for advanced suite variables or quality assurance samples
- Samples taken for laboratory analysis
- Whether any samples were not collected, and reason
- Water level
- Any other required metadata for NPSTORET or Aquarius data entry
- Time of departure

All entries should be made clearly. If an incorrect entry is made, a single heavy line should be drawn through the incorrect entry and the correction made. All corrections should be initialed and dated. The completed field forms will be maintained in chronological order by station, copied into site binders and the originals maintained on file indefinitely. Field data are reviewed annually by network personnel (see SOP #8, Data Entry and Management, for details).

12.7.2 Site/Sample Identity Codes

Sampling stations will be identified by park and water body according to Network guidance. Information on the sampling station and park will comply with NPSTORET and Aquarius requirements.

12.7.3 Data Transfer

Import field and laboratory data into NPSTORET or Aquarius as soon as possible after receiving the data, according to SOP #8. Field forms are the only written records of field measurements; place copies in office binders and keep originals on file indefinitely. Program sampling data and associated records are archived and stored in the GLKN Ashland Office. Boxes are numbered consecutively by year, project, and station number.

Personnel importing data into the database should take care to ensure that units are consistent for laboratory data. Different laboratories may report analytical results in different units (mg/L vs. µg/L) for the same analyte. Follow SOP #8, Data Entry and Management, when importing data.

Contract laboratories should use the STORET-supported laboratory remark codes (Table 12-7) or provide a map relating their remark codes to these. Unlisted remark codes should be discussed for possible addition by the USEPA to the STORET codes. The detection descriptors to be used in data entry to STORET are listed in Table 12-8.

Table 12-7. STORET-supported laboratory remark codes.

Code	Definition
AL	Aldol condensation present. Analyte may not be present.
CNT	Non-acceptable colony counts.
EHT	Sample or extract held beyond acceptable holding time.
FBK	Analyte found in blank. Sample contamination indicated.
FDB	Failed. Dry blank not acceptable.
FDC	Failed. Drift check not acceptable.
FIS	Failed. Internal standard not acceptable.
FLD	Failed. Lab duplicate not acceptable.
FFD	Failed. Field duplicate not acceptable.
FFB	Failed. Field blank not acceptable.

Table 12-7 (continued). STORET-supported laboratory remark codes.

Code	Definition
FFS	Failed. Field spike not acceptable.
FFT	Failed. Trip blank not acceptable.
FLC	Failed. Linearity check did not meet quality criterion.
FLS	Failed. Lab spike recovery not acceptable.
FMS	Failed. Matrix spike recovery not acceptable.
FPC	Failed. Lab performance check not acceptable.
FQC	Failed. Quality control criteria exceeded during analysis.
FRS	Failed. Internal reference sample not acceptable.
FSP	Failed. Surrogate spike recovery not acceptable.
FSB	Failed. Spiked field blank recovery not acceptable.
FSL	Failed. Spiked lab blank recovery not acceptable.
INT	Interference suspected. Analyte may not be present.
ISP	Improper sample preservation noted. Analysis performed.
LIS	Lab internal standard(s) added to sample.
LLS	Value less than lower quality control standard.
PRE	Presumptive evidence that analyte is present.
NJ	TIC, Tentatively Identified Compound, result is approximate
N	TIC, Tentatively Identified Compound, presumptive id only
OUT	Result value is defined as an outlier by data owner
SUS	Result value is defined as suspect by data owner

Table 12-8. STORET detection descriptors.

Detected and Quantified
*Non-detect
*Present >QL
*Present <QL
*Not Reported
*Present

12.7.4 Sample Transfer

Conduct sample processing per SOP #7. Refrigerate or freeze samples, as required. Conduct in-house laboratory work and package samples for sending to contract analytical laboratory. Sample preservation and conditions for shipment will differ for each parameter; these needs should be discussed ahead of time with the contract laboratory and documented. Fill out the analytical request form and chain-of-custody (COC) form provided by the laboratory (examples are included in SOP #7). The COC form is used to document the taking, shipment, and receipt of samples. The laboratory will use the COC to check samples into the analytical process. Clean all transfer bottles and equipment according to SOP #7. Water samples are maintained as close to sampling conditions as possible by shipping on ice. Chain-of-custody documentation will be maintained. A chain-of-custody includes not only the form, but all references to the sample in any form, document, or log book that allow tracing the sample back to its collection, and documents the possession of the samples from the time they were collected until the sample analytical results are received.

12.8 Analytical Methods

Field equipment and contract laboratories are likely to change during the course of this long-term monitoring project. Documentation will be maintained, therefore, as regards each multiprobe and contract laboratory's ability to meet the data quality objectives of this project. Forms for this purpose are included in Attachment A. These completed forms for each multiprobe and contract laboratory will be maintained along with the maintenance logs for the multiprobe and QAPPs for each analytical laboratory.

12.8.1 Field Methods

One important aspect in the low uncertainty/bias and high precision of a water quality monitoring program is the correct selection of probes for measuring field variable and their subsequent calibration and maintenance. Table 12-3 lists typical field sensor performance specifications that should be expected from monitoring equipment for this protocol. Table 12-9 summarizes the ideal calibration frequency and minimum acceptance criteria for these sensor probes. The reality of logistical constraints at back country sites may preclude calibration and checks of calibration at the ideal frequency.

Table 12-9. Ideal calibration frequencies and acceptance criteria for field instruments.

Parameter	USEPA Method	Minimum Calibration Frequency and QC checks	Acceptance Criteria	Corrective Actions
Temperature	170.1	Annually, 2-point check with NIST thermometer	$\pm 1.0^{\circ}\text{C}$	Re-test with a different thermometer; repeat measurement
Specific Conductance (SC25)	120.1	Daily, prior to field mobilization; calibration check prior to each round of sampling; 10% of the readings taken each day must be duplicated or a minimum of 1 reading if fewer than 10 samples are read.	$\pm 5\%$ RPD 10%	Re-test; check low battery indicator; use a different meter; use different standards; repeat measurement

Table 12-9 (continued). Ideal calibration frequencies and acceptance criteria for field instruments.

Parameter	USEPA Method	Minimum Calibration Frequency and QC checks	Acceptance Criteria	Corrective Actions
pH	150.1	Daily, prior to field mobilization (two buffers should be selected that bracket the anticipated pH of the water body to be sampled)	±0.05 pH unit	Re-test; check low battery indicator; use different standards; repeat measurement; don't move cords or cause friction/static
		Calibration check w/ third buffer prior to each round of sampling; check with low ionic strength buffer in addition, if conductivity is <50 µS/cm	±0.1 pH unit	Re-test; check low battery indicator; use different standards; repeat measurement; don't move cords or cause friction/static
		10% of the readings taken each day must be duplicated or a minimum of 1 reading if fewer than 10 samples are read.	RPD 10%	Re-test; check low battery indicator; use different standards; repeat measurement; don't move cords or cause friction/static
Dissolved Oxygen	360.1	Daily, prior to field mobilization; check at the field site if elevation or barometric pressure changed since calibration	0.2 mg/L concentration or ±10% saturation	Re-enter altitude; re-test; check low battery indicator; check membrane for wrinkles, tears or air bubbles; replace membrane; use a different meter; repeat measurement; allow more time for stabilization
Depth	--	Daily, prior to field mobilization, check at the field site. Check annually against commercially purchased brass sash chain labeled every 0.5 m to ensure that it reads zero at the surface and varies <0.3 m for depths <10 m and no more than 2% for greater depths.	±0.1 m	Retest, check low battery indicator; repeat measurement; use with accurately calibrated line
Transparency tube	--	Transparency tubes have a 100 or 120 cm scale; ensure tube is clean	±1.0 cm for transparency tube	--
Marked lines (e.g., Secchi, Van Dorn)	--	Check markings annually against brass sash chain. If lines are heated (for decontamination) check prior to each round of sampling.	±1%, 0–10 m ±2%, >10 m	Re-mark line.

Ensure calibration standards are not used beyond expiration dates. Refer to SOP #6 for an explanation of each parameter measured, guidelines on potential field measurement problems, and other details on performing calibration checks. Keep the manufacturer's manual with each instrument for aid in troubleshooting.

Used calibration solutions, in general, may be rinsed down a sink with water after consideration of the wastewater treatment system available to that sink. Safety data sheets (SDS) which are sent with manufacturer purchased calibration solutions should be kept on file. These documents describe the flammability, toxicity, and other safety hazards of reagents. Some reagents may include constituents toxic to aquatic life. These should not be rinsed down a sink in any large quantities in primitive areas where the ultimate destination of wastewater is the aquatic environment. These reagents should be collected in a leak-proof container that is properly marked, and disposed of in an adequate treatment system.

Calibration logs for multi-parameter sondes will be maintained and will document the frequency of calibration and calibration checks. Data reporting units will be standardized by using the data sheet. Keep calibration and maintenance logs for multiparameter sondes with the sonde during the sampling season. Logs will later be archived at the Network office in Ashland, Wisconsin. A new log will be started for each field season.

12.8.2 Field Equipment Performance Criteria

Estimates of the completeness, representativeness, precision, comparability, and systematic error of data generated by the use of a multiprobe will be estimated and documented according to this SOP.

Data will be considered representative of the site when procedures detailed in the large rivers protocol narrative and associated SOPs are followed. Actual sampling location and changes due to field conditions will be documented on the field forms. Data will be comparable year-to-year as objectives and methods chosen are consistent within documented NPS GLKN sampling procedures.

12.8.2.1 Field Data Completeness

The completeness of data collected, or percentage of intended field measurements that were actually made, will be calculated at year's end according to the formula below.

$$\text{Percent completeness} = \left(\frac{\# \text{ samples_collected}}{\# \text{ samples_planned}} \right) \times 100$$

12.8.2.2 Instrument Sensitivity

Sensitivity of the multiprobe will be documented through an estimation of the limits of detection known as alternative measurement sensitivity (AMS). The AMS for each multiprobe will be estimated annually, and for any new multiprobes, by taking multiple (at least seven) measurements in a field simulation for each parameter. Alternative Measurement Sensitivity (AMS) is a two-sided estimation based on the 99% confidence interval for sample size seven. Of interest is the size of the difference between two individual values that can be considered to actually be a true difference. This calculation differs from the MDL, which is a one-sided true difference from zero. The standard deviation will then be calculated and used in the estimation of AMS ($3.708 \times \text{SD}$). This estimation will be carried out and documentation kept for a new probe and annually thereafter (Attachment A). Documentation will be kept in the project files.

12.8.2.3 Instrument Precision

The precision of the multiprobe will be estimated through the use of duplicates. Duplicates will be measured each day in the field at a rate of 10% (or 1 per park sampled if fewer than 10 sites are sampled). A reading will be considered a duplicate when it is repeated at a particular sampling site immediately after it is initially taken. Only the surface reading need be replicated. This will reduce the contribution of variability from the variability of the site itself. Precision calculations are made by way of a Relative Percent Difference (RPD). Measurement Quality Objective (MQO) for precision: The RPD should be less than or equal to 10%; details and corrective actions are listed in Table 12-10. Calculate precision as follows:

$$Precision = \left(\frac{A - B}{(A + B) / 2} \right) \times 100$$

Where A is the first reading, and B is the second reading taken immediately after the first.

Table 12-10. Frequency, acceptable range, and corrective actions for quality control (QC) samples.

Type of QC Sample	Frequency	Acceptable Range for MQOs	Corrective Action
Field Duplicates (samples)	Minimum of 1 per trip per parameter or 10% of all samples per parameter per day	All parameters $\pm 10\%$ RPD, chlorophyll-a, TSS and nutrients $\pm 30\%$	Audit field personnel and verify sample collection procedure; resample; reanalyze; revise SOP; audit and train field personnel; project manager determines whether associated data is usable
Field Duplicates (multi-probes)	Minimum of 1 per trip per parameter or 10% of all samples per parameter per day	All parameters $\pm 10\%$ RPD	Re-calibrate instrument; replace batteries; perform instrument field check with different standards; repair or replace instrument; notify management; audit and train field personnel; project manager determines whether
Laboratory Control QC (bias)	One each per analytical batch, minimum 1 per 20 samples	All parameters $\pm 15\%$ RPD	According to laboratory QAPP, ensure re-calibration, re-analysis, and documentation.

12.8.2.4 Instrument Bias

The systematic error/bias of the multiprobe will be estimated through the use of reference solutions. Reference solutions will be measured prior to each round of sampling, in the lab or field. The results will be documented and used as a calibration check according to Table 12-9 as well as for a long-term estimate of systematic error/bias. Bias calculations are made by way of a Percent Difference (PD). Calculate bias as follows:

$$Bias = \left(\frac{Y - X}{X} \right) \times 100$$

Where X is the known (or expected) amount and Y is the measured concentration.

12.8.3 Laboratory

The ability to use environmental data to reveal long-term trends requires consistent analytical methods and detection limits. The NPS recommendation is to use only laboratories with NELAC certification or at least certification by the state programs that also use the laboratory. The chosen contract laboratories must prove their capability annually through participation in blind quality control checks and other methods prescribed by the states in which they receive certification and/or federal programs in which they participate. Copies of certifications for each analyte and/or method will be kept on file along with Quality Assurance Project Plans (QAPPs) for each laboratory contracted for the duration of this monitoring effort.

The method used in calculating method detection limits (MDLs) and method limit of quantitation (ML) or laboratory reporting limits (Lt-RL for USGS labs) may differ for each contract laboratory. Most laboratories routinely recalculate MDLs, MLs, and QC sample control limits using repeated measurement of standard samples or multiple percent recoveries on a quarterly or annual basis. The GLKN will request and maintain copies of this information as most recently calculated for the relevant analytical methods to assist in the selection of contract laboratories, data validation, and AMS calculations.

A checklist to be used in selecting contract laboratories and documenting their compliance with GLKN QA/QC expectations is included in Attachment A.

Great Lakes Network staff validating laboratory data for database entry should take care to ensure that data are entered in consistent units, as different laboratories may report results in different units (mg/L or µg/L) for the same analytes. A checklist to be used in data validation is included in Attachment B.

12.8.4 Laboratory Performance Criteria

Comparability, representativeness, precision, systematic error, and completeness of data generated by contract laboratories will be estimated and documented according to this SOP during data validation. Table 12-10 can be used as a guide. Unless otherwise justified (for example, to be consistent with state requirements), in no case will Measurement Quality Objectives less stringent than the following be accepted:

- Precision—A maximum of 10% RPD for all lab parameters except chlorophyll-*a*, nutrients, and total suspended solids, for which the maximum RPD is 30%.
- Bias—A maximum of 15% PD for all parameters, or state credible data defaults, whichever is more stringent.

All MQOs will be used as data rejection criteria.

The data delivery package must contain QC sample results and an explanation of new STORET and NPSTORET compatible laboratory flags. A checklist to be used for documenting this procedure is included in Attachment B.

Data generated will be considered representative of the particular site when samples are taken according to the sampling protocol for each sampling objective. Actual sampling location and changes due to field conditions will be documented on the field forms. Data will be comparable year-to-year as objectives and methods chosen are consistent within documented sampling procedures.

12.8.4.1 Laboratory Data Completeness

The completeness of data analyzed, or the percentage of intended sample measurements that were actually made, will be calculated at year's end according to the formula below. Data points may be missed due to site conditions, sample container breakage, or disqualified analyses due to control limit exceedances in the laboratory. The reasons for missing data should be documented.

$$\text{Percent completeness} = \left(\frac{\# \text{samples_analyzed}}{\# \text{samples_planned}} \right) \times 100$$

Determining required sample sizes and attendant completeness goals was done in a stepwise manner, considering desired statistical power and minimum detectable differences (see discussions in the protocol narrative). Should percent completeness ever fall below sample sizes needed for analysis of trends, adaptive changes will be made to ensure it does not happen again.

12.8.4.2 Laboratory Sensitivity

Measurement sensitivity is estimated in laboratory analysis through the use of signal to noise ratios or the standard deviation of repeat measurements of a low level reference standard. The method each laboratory uses to calculate MDLs and MLs will be documented using the data validation check list. This information will be reviewed during data validation and kept along with other QA/QC information for each laboratory.

12.8.4.3 Laboratory Precision

Measurement precision and bias are estimated using a number of QC samples during a round of sample analysis for each method. A reference standard of a certified known concentration is analyzed along with samples at a rate specified by the laboratory's QA/QC program. The results of these samples are reported and the percent difference calculated. Laboratory duplicates of field samples are also analyzed. This is a field sample that is split or subject to repeat analysis within the laboratory to estimate and control precision repeatability. Relative percent differences are calculated for laboratory duplicates. The percent difference and RPD of these QC samples should fall within the control limits specified by the laboratory (usually 10% to 20% depending on the type of analysis made).

Field duplicates are to be sent to the analytical laboratory at a rate of at least one in every 10 samples. A field duplicate is a split sample, or at minimum, a sample co-located in time and space. The duplicate is usually given a separate sample identity on the sample label and forms, and it is noted on the field form for which sample this is a duplicate. This is a separate check on precision and repeatability of the laboratory analysis. The precision measurement is calculated using the Relative Percent Difference (RPD) between duplicate sample results per analyte. Precision estimates should be performed within seven days of receipt of laboratory results as part of data validation. Acceptance criteria and corrective actions are summarized in Table 12-10.

12.8.4.4 Laboratory Bias

Systematic error/bias (formerly referred to as percent recovery) is estimated for each laboratory method through the analysis of spiked samples and/or certified reference materials. Spiked sample analyses are conducted by the contract laboratory to ensure the reported data are accurate, or compare favorably to the true values. A spiked sample consists of a sample with a known concentration of analyte added before any sample preparation procedures are carried out. Sample and spiked sample are analyzed using normal procedures and the percent spike difference is calculated. Percent difference is calculated as bias (see equation in Section 12.8.2.4).

Other sources of systematic error in monitoring programs include sample cross contamination from field sampling equipment used to handle a multiple number of samples. Equipment blanks are used to estimate whether systematic error is added to sample data during sample handling. The integrated sampler is to be rinsed in the field three times at each site prior to taking samples. Compositing jugs that are not site-dedicated may also be a source of cross contamination. Collect an equipment blank periodically, as follows:

1. In between sample sites, rinse the equipment used to transfer water samples (integrating sampler or compositing jug) with laboratory reagent-grade water three times and discard.
2. Rinse with a fourth aliquot and save this aliquot to lab bottles as if it were a river water sample to be analyzed.

This sample is labeled equipment blank and information kept on a datasheet describing the source of the blank. Results for all parameters should be non-detect. This type of blank is a check for cross contamination between sampling sites and control for bias introduced by cross contamination.

The laboratory QAPP should define control limits to be used during analysis of samples. If the analysis QC samples are not meeting the control limits, the analysis is usually repeated after re-calibration. This protocol should be documented in the lab's QAPP. During data validation GLKN will ensure that the laboratory performance meets the precision and bias MQOs tabulated in Table 12-10.

12.8.5 Changing Methods and Documenting Cumulative Bias

When a field method is changed, the cumulative bias, or change in sampling results due to the method change alone, should be estimated. Cumulative bias can become significant over time even though changes in methods are small. When change occurs due to a scheduled change in staff, both the new staff member and old staff member should perform side-by-side field measurements several times (minimum of seven) during training when possible. The results for both will be compared as below.

When purchasing a new probe, it should be used for a minimum of seven measurements side-by-side with the existing probe when possible.

When a change in laboratory methods is made by the laboratory, the laboratory will be responsible for estimating any bias introduced by the change. When a laboratory contract is changed to a

different laboratory, a minimum of seven samples will be split for analysis in both labs. Data from these comparisons will be used to calculate a percent difference (PD) and fraction of change:

$$PD = \left(\frac{new - old}{old} \right) \times 100$$

$$Fraction_of_change = \left(\frac{old}{new} \right)$$

Where *old* = data from the older or original method and *new* = data from the new or replacement method.

Ideally, the average PD (two-sided 95% t-distribution confidence interval) will be within 20% of the mean of the old or the new method measurements. If this is not the case, the number of overlapping measures should be increased until this criterion is met (Irwin 2006). New values can then be normalized to the old method value by multiplying them by the fraction of change.

Any bias should be stated clearly and documented. Documentation will be stored within each site file and should also include:

- all raw data pairs for future use
- standard deviation
- the MDL or AMS of the method
- number of paired samples
- 95% t-distribution confidence interval for the measurement difference
- dates between which overlapping measurements were made
- date documentation calculation was made

These comparison of methods will be carried out, when possible, in order to document any bias that is introduced into the long-term data by a change in methodology and not in the actual environment. Due to the logistical realities imposed by sampling in remote locations and the reality of equipment loss or damage and abrupt changes in staff, this type of comparison may not always be possible.

12.9 Instrument Calibration Frequency, Inspection, and Maintenance

Instruments used in field measurement or laboratory analysis often require frequent inspection and maintenance to ensure they are in good working order. Calibration frequencies differ by instrument and should be spelled out in SOPs for both field and laboratory work to ensure consistency.

12.9.1 Field Instruments

Each instrument must have a logbook. The calibration schedule must be observed, using fresh calibration standards. Calibration solutions may be disposed of by rinsing down the work space sink if the contents do not include constituents which would harm the pipes or aquatic life on the

receiving end of the available wastewater treatment (see relevant SDS sheets). Calibration checks must be documented.

When sensor probes are to be stored for extended periods of time, thoroughly clean sensors, remove batteries, and store sonde according to specific instructions in SOP #11 and manufacturer's manual. Store calibration standards and electrolyte solutions in a temperature-controlled environment.

12.9.2 Laboratory Instruments

Instrument calibration and verification is performed at least once a day for each analytical method prior to the analysis of samples. Each laboratory specifies its own procedure, which should include multiple point calibrations plus a blank, continuing calibration verification, and final calibration verification at the end of an analytical run. Each laboratory should define the procedures in a QAPP. Review of these procedures will be documented using the laboratory checklist in Attachment B.

12.10 Inspection and Acceptance of Supplies and Consumables

Monitoring water quality requires many pieces of equipment and a large amount of supplies. Table 12-11 lists the SOPs that include checklists of supplies and equipment. When new equipment is received or equipment is returned after repair, inspect it for flaws and test it to ensure proper functioning. When supplies are received, inspect them to ensure containers are properly sealed. Ensure reagent containers are dated upon receipt and upon opening. Commercially purchased calibration standards come with an expiration date that must be observed.

Table 12-11. Checklists of equipment and supplies for monitoring water quality of large rivers.

Checklist	Location
Safety equipment and supplies	SOP #2
Decontamination equipment and supplies	SOP #5
Field equipment and supplies	SOP #6
Laboratory equipment and supplies	SOP #7

12.11 Records Management

All records must be kept, from field activity through sample results. Required metadata will be complete. Table 12-12 summarizes the QA/QC procedures related to data management. For more details on the Great Lakes Network's overall strategy for metadata generation, management, and distribution see chapter 8, Data Documentation, of GLKN's Data Management Plan (Hart and Gafvert 2005) and the appendices of that document.

Table 12-12. Summary of QA/QC procedures pertaining to data management.

Procedure	Description
Instrument calibration logs	Each instrument must have a logbook.
Field forms	Field forms are the only written record of field measurements. Place copies in site binders and keep originals on file indefinitely.
Estimating precision	The precision measurement is calculated using the Relative Percent Difference (RPD) between duplicate sample results per analyte. Precision estimates should be performed within 7 days of receipt of laboratory results.
Electronic data entry	Approximately 10% of electronic data entries should be spot checked for errors on a random basis. If errors are found, another 10% are spot checked.
Data archiving	Sampling data and associated records are archived in boxes and stored at the GLKN Ashland office. Boxes are numbered consecutively by year, project, and station number.
Data validation	Data validation is the process that determines whether quality control objectives for data collection were met.
Data validation reports	Data validation reports provide a narrative that discusses any deviations from QA/QC procedures and the impacts of those deviations.
Data verification	Data verification demonstrates that a data set will qualify as credible data.
Data verification reports	Data verification reports document the results of the data verification procedure.
Data qualification codes	Data must be fully qualified before uploading to the Water Resources Division's NPSTORET

Data will be archived according to SOP #8 (Data Entry and Management) for digital data, paper copies, and field forms. Field forms are maintained indefinitely. Brief characterizations of the data from each NPS unit that was sampled and the Network as a whole will be performed each year, after all QA/QC procedures have been completed. For each station sampled, these descriptive statistics will include mean, median, maximum and minimum values, skew, kurtosis, and measures of variability (e.g., CV, standard error, variance) for each water quality variable. Understanding of landscape-scale dynamics will be provided by analyses of variability among stations within the domain of interest (e.g., park unit), following one of several approaches described in SOP #9, Data Analysis.

A 'Memo-to-file' will be used to document any decisions or corrections that are made. This memo will include the date, name of author, site or sample referred to, a description of the problem or error and a statement describing the decision made or action taken. The memo will be archived with the appropriate site data and files.

12.12 Assessment and Oversight

It is the project manager's responsibility to make sure each component of this and other SOPs are followed.

12.12.1 Corrective Responsive Actions

The project manager, in consultation with experienced professionals, will be responsible for taking corrective responsive action in the case in which QA/QC is not followed or in the case of an unexpected event. Responsive action is often needed in the event of broken sample bottles, missing data, errors on field sheets, changes due to field conditions, problematic analyses, and other events that do not fall within the standard operating procedures. A “Memo-to-file” will be used to document any decisions or corrections that are made. This memo will include the date, name of author, site or sample referred to, a description of the problem or error and a statement describing the decision made or action taken. The memo will be archived with the appropriate site data and files.

12.13 Reports to Management

Routine data summaries will occur annually for large rivers. Summary reports will be produced periodically. Web-based visualization and data summary tools for large river water quality data are under development. These tools and summary reports will be provided to park resource managers, park partners, and the public.

More comprehensive analyses of trends will occur for most parameters after three or more years of sampling. For stations that are located where no previous monitoring has occurred, this period of time tends to be the minimum needed to establish a time series sufficiently powerful to detect meaningful levels of change (e.g., 20%) through time. The target audience of the analysis and synthesis reports will be the parks, the Network, both regional and Service-wide I&M, and the broader scientific community. Drafts will be reviewed internally and sent to the parks, and possibly outside sources, for further review. The extent of review will depend on how analytically complicated the methods are and the gravity of inference and recommendations.

12.14 Data Validation, Certification, and Usability

Data validation is the method in which data are proven or disproved to be accurate. This process involves the review of the results of all measurements, samples, and QC samples. Field data sheets and laboratory data are reviewed for transcription errors, completeness, verification of calibration and quality control check samples or standards.

A checklist for use in data validation is presented in Attachment B. Data are validated by comparing the actual estimates for MDL and ML, precision, bias/uncertainty for each analytical run to the expected quality of results. Once this information is documented, the decision is made by the project manager on the usability of the data. Data may be flagged as between MDL and ML and therefore highly uncertain (J). Data may be flagged (B) if the analyte was detected in the blank at a concentration similar to that in the sample (laboratory flagging rules may differ slightly). These types of flagged data contain much more uncertainty than unflagged data. The decision must be made whether or not to add them to the database or merely maintain them in files for future reference. Some investigation into the sampling method or laboratory method may be needed if ‘B’ flagged data are a continual occurrence or if the required MLs are repeatedly unmet.

NPSTORET itself includes some form-based QA/QC tests on data entry. The system does not allow entry of results below the ML but will flag the data as “present, not quantified”. The project manager, in consultation with other NPS or outside experts, will flag data when they fall outside of expected limits. Laboratory results below detection limits (ND) will be kept in the archives as ‘ND’ and will be handled statistically according to recent literature (e.g., Helsel 2005) in summary and analysis reports. Aquarius provides QAQC tools to check for single value or rate of change data outliers, for example.

Once returned to GLKN by WRD, and after appropriate documentation is completed, the NPSTORET datasets are officially considered certified. A process for certifying data imported into Aquarius is being developed by WRD. Data certification is a benchmark in the project information management process that indicates that: 1) the data are complete for the period of record; 2) they have undergone and passed the quality assurance checks; and 3) that they are appropriately documented and in a condition for archiving, posting and distribution as appropriate.

Certification is not intended to imply that the data are completely free of errors or inconsistencies, which may or may not have been detected during quality assurance reviews. Only qualified users who have been trained and given edit permissions are allowed to edit data in NPSTORET or Aquarius. These procedures protect the integrity of the data and allow the history of each data record to be traced.

12.15 Reconciliation with Data Quality Objectives

Once data validation is complete, the project manager then calculates percent completeness, compares the results to the data quality objectives, and determines whether or not the data meet the objectives for both the field and laboratory components of the project.

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Attachment 12A. Checklists for Selection of Multiprobe and Selection of Laboratory.

Multiprobe Checklist

Probe _____ Purchased _____

Serial Number _____ Location used _____

Manufacturer's Stated Limits

Sensor	Range	Reporting Resolution	Bias	Stabilization Criteria
Temperature				
Sp. Cond. (SC25)				
pH				
DO (Conc.)				
DO (% sat.)				
Depth – Z (pressure sensor)				

Alternative measurement sensitivity by multiple field readings

Location _____ Performed by _____

Date _____ Began calibration log _____

Sensor	1	2	3	4	5	6	7
Temperature							
Specific Conductivity (SC25)							
pH (sensor units)							
DO (Conc.)							
DO (% sat.)							
Depth – Z (pressure sensor)							

Calculations

For each parameter calculate the standard deviation of the 7 readings.

Sensor	Standard deviation	AMS 3.708x sd		
Temperature				
Specific Conductivity (SC25)				
pH (sensor units)				
DO (Conc.)				
DO (% sat.)				
Depth – Z (pressure sensor)				

Alternate Measurement Sensitivity is a two-sided estimation based on the 99% confidence interval for sample size 7. Of interest is the size of the difference between two individual values that can be considered to actually be a true difference. The MDL is a one-sided true difference from zero.

Contract Laboratory Checklist

Laboratory:

Address:

Contact person:

_____ Received QAPP

_____ Received copy of certifications

_____ Received a list of analytical methods used

_____ Define the limits of detection and limit of quantitation calculation method
(can lab report as MDL and ML if currently calculated as LOD and LOQ?)

_____ Received a copy of latest MDL/ML/control limits calculations for relevant methods

_____ MDL/ML as listed meet project needs
(list any analyte for which the ML requirement is not met – discuss options with lab)

_____ Received successful interlaboratory participation documentation

_____ Sample handling log in and COC are documented in the QAPP

_____ Equipment maintenance and calibration procedures are documented in the QAPP

_____ Internal QA/QC documented in QAPP

Control limits calculations are made by what method
(QC includes blanks, duplicates, spikes, reference standards and LCS)

Calibration curves cover level of analytical interest
(QC includes ICV and ICB, and CCV and CCB)

Reporting data flags used include

Attach all copies

Reviewed by _____

Date _____

Attachment 12B. Checklists for data validation.

Field Measurement Validation Checklist

(one sheet per Park sampled)

Sampling Unit _____

Date Sampled _____

Date Reviewed _____

Reviewed by _____

_____ All field forms have been received (data sheet, flow, etc.)

_____ Multiprobe was calibrated correctly

_____ Multiprobe post-calibration checks were successful

_____ Field duplicates were within range

_____ Multiprobe end of use calibration checks were successful

_____ There were no obvious trends in data taken from any sensor during the sampling day

_____ Equipment blanks sent to lab included

Laboratory Data Validation Checklist

Sample set from _____

Date taken _____

Received data _____

Reviewed by _____

Review date _____

_____ Samples received by lab at proper temperature (look at COC copy from lab)

_____ Holding time limits met

_____ Analytical methods used in analyses were those agreed upon

_____ Useable MDL and ML achieved in this analytical run

_____ Calibration procedures were followed

QC samples control limits applicable to this analytical batch

Lab blank _____

Lab dup _____

Lab LCS _____

Lab spikes _____

Field dup _____

Eq blank _____

CCVs _____

QC samples within range expected

Lab blank _____

Lab dup _____

Lab LCS _____

Lab spikes _____

Field dup _____

Eq blank _____

CCVs _____

Lab notes or flags

Samples rejected

Signature of Reviewer _____

Standard Operating Procedure #13: Procedure for Revising the Protocol

Version 1.1

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Contents of SOP 13

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Revision History Log

The following table lists all edits and amendments to this document since the original publication date. Information entered in the log must be complete and concise. Users of this standard operating procedure will promptly notify the project manager and/or the Great Lakes Network (GLKN) data manager about recommended and required changes. The project manager must review and incorporate all changes, complete the revision history log, and change the date and version number on the title page and in the header of the document file. For complete instructions, please refer to Revising the Protocol, SOP #13.

Previous Version #	Revision Date	Author (with title and affiliation)	Location in Document and Concise Description of Revision	Reason for Change	New Version #
1.0	3/18/16	David VanderMeulen, GLKN	Updated the NPS authorship and contact author, added information about obtaining a new document number, noted that data manager will repost new version of narrative to IRMA if major changes are made, fixed a handful of grammatical errors throughout	To clarify and update SOP based on changes that have occurred since Version 1.0 was published in 2008	1.1

*Add rows as needed for each change or set of changes tied to an updated version number

13.0 Introduction

Because of the long-term nature of the National Park Service's (NPS) monitoring program, the projects must necessarily accommodate change. Refined field methods, advances in analysis techniques, and feedback from field crews and project managers can all contribute to improving the monitoring protocol. The purpose of the current SOP is to define a systematic and routine process for incorporating these changes into the protocol.

13.1 Steps for Revising the Protocol

1. Attempt to incorporate the changes by first modifying only the SOP(s), without making changes to the protocol narrative. However, if it is clear that changes will also be needed on the narrative, then revise it as well.
2. Make all revisions using the Track Changes feature of Microsoft Word. For minor changes, at least one other person must review the revision. If the change is more extensive, a discussion of the changes by Network staff is warranted before acceptance of the revision. For major changes, review from outside of the Network should be sought. Examples of major changes include modifications of the sampling design, significantly altered field methods, and revised analysis techniques.
3. Record the changes in the revision history log of the SOP and/or in the narrative, as appropriate. Include the date of revision, full name(s) and affiliation(s) of author(s), description of and reasons for the changes, and section of SOP or narrative where changes were made.
4. Rename the version of the SOP and/or narrative. For minor changes, only revise the version number after the decimal point (e.g., change V. 1.1 to V. 1.2). For major changes, revise the number before the decimal point (e.g., V. 2.3 to 3.0) and obtain a new Natural Resources Report number. Also change the version number of the SOP or protocol in the header or footer, as appropriate.
5. Notify the data manager of the change(s) so that the metadata of the project database will be updated. If major changes are made to the protocol narrative the data manager will also repost the updated narrative to the NPS Integration of Natural Resource Applications website.
6. Distribute the revised version to all appropriate parties, including the members of the field crew and appropriate GLKN staff. The revised version must also be posted on the Network's website.
7. Maintain a library of previous versions. Such historical information may be crucial for understanding, interpreting, and analyzing data.

The Department of the Interior protects and manages the nation's natural resources and cultural heritage; provides scientific and other information about those resources; and honors its special responsibilities to American Indians, Alaska Natives, and affiliated Island Communities.

NPS 920/133725, August 2016

National Park Service
U.S. Department of the Interior



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