

Quality Assurance Project Plan

Hoonah BEACH Program

April 2022



Prepared by:

**Nonpoint Source Pollution Program
Division of Water
Alaska Department of Environmental Conservation
State of Alaska**

A. Project Management Elements

A.1 Title and Approvals

Title: Quality Assurance Project Plan for 2022 Hoonah BEACH Program

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A.2 Distribution List

This list includes the names and addresses of those who receive copies of the approved QAPP and subsequent revisions.

<i>Table 1: Distribution List</i>				
NAME	POSITION	AGENCY/ Company	DIVISION/BRANCH/ SECTION	CONTACT INFORMATION
Laura Eldred	NPS Section Manger	DEC	Division of Water/ WQ / Non-Point Source	907-376-1855 laura.eldred@alaska.gov
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A.3 Project Task/Organization

Dues and responsibilities of key individuals are listed below:

A.3.1 DEC Staff:

- **NPS Section Manager** – Responsible for overall technical and contractual management of the project.
- **DEC QA/QC Officer** – Responsible for QA review and approval of plan and oversight of QA activities ensuring collected data meets project’s stated data quality goals. Conducts field audits, data audits, QA review of blind lab performance, evaluation of samples, and lab audits.
- **Project Manager/ Field Coordinator** – Responsible for overall technical and contractual management of the project. If DEC staff have direct responsibility for sample collection and analysis of data results, the DEC Project Manager/s assume the responsibilities of the Lead Field Sampler/Project Manager.
- **Laboratory Manager** - Responsible for the overall review and approval of contracted laboratory analytical work, responding to sample result inquiries and method specific details. Responsible for QA/QC of laboratory analysis as specified in the QAPP and reviews and verifies the validity of sample data results as specified in the QAPP and appropriate EPA approved analytical methods. These duties may be assigned to a lab appointed project manager.

A.3.2 Southeast Alaska Watershed Coalition Staff

- **Project Manager/Project QA Officer** – Responsible for overall technical and contractual management of the project. If SAWC staff have direct responsibility for sample collection and analysis of data results, the SAWC Project Manager assumes the responsibilities of the Lead Field Sampler/Project Manager. Responsible to ensure all monitoring complies with the QAPP specified criteria. This is accomplished through routine technical assessments of the sample collection, analysis, and data reporting process. Assessments may include but are not limited to activities such as: on-site field audits, data audits, QA review of blind lab performance evaluation samples, and lab audits. These assessments are performed independent of overall project management.
- **Laboratory Manager** – Responsible for the overall review and approval of contracted laboratory analytical work, responding to sample result inquiries and method specific details. Responsible for QA/QC of laboratory analysis as specified in the QAPP and reviews and verifies the validity of sample data results as specified in the QAPP and appropriate EPA approved analytical methods.

A.3.3 HIA Staff:

- **Lead Field Sampler** – Responsible for sampling preparation, sample collection, sample preservation, transportation of samples to laboratory for analysis, receipt of data and transmittal of data to Project Manager. The individual will procure personal equipment of field personnel, coordinate with laboratories in planning sampling equipment needs, obtain supplies for and prepare daily sampling kits prior to departure for field location, travel to the field location, prepare necessary preservatives while in the field, perform site reconnaissance, collect

site specific parameters, collect water samples, prepare samples for shipping, transport samples to laboratory, alert laboratory of successful sampling event, receive data from laboratory, verify sample result data is reliable and submit the data and all applicable QA/QC results to the SAWC and DEC Project Managers.

- **Field Support Personnel** - Responsible for accompanying Lead Field Sampler into the field and supporting Lead Field Sampler during sampling. The individual will travel with the Lead Field Sampler to the field location, accompany the Lead Field Sampler to sampling sites, and support Lead Field Sampler in sampling tasks.

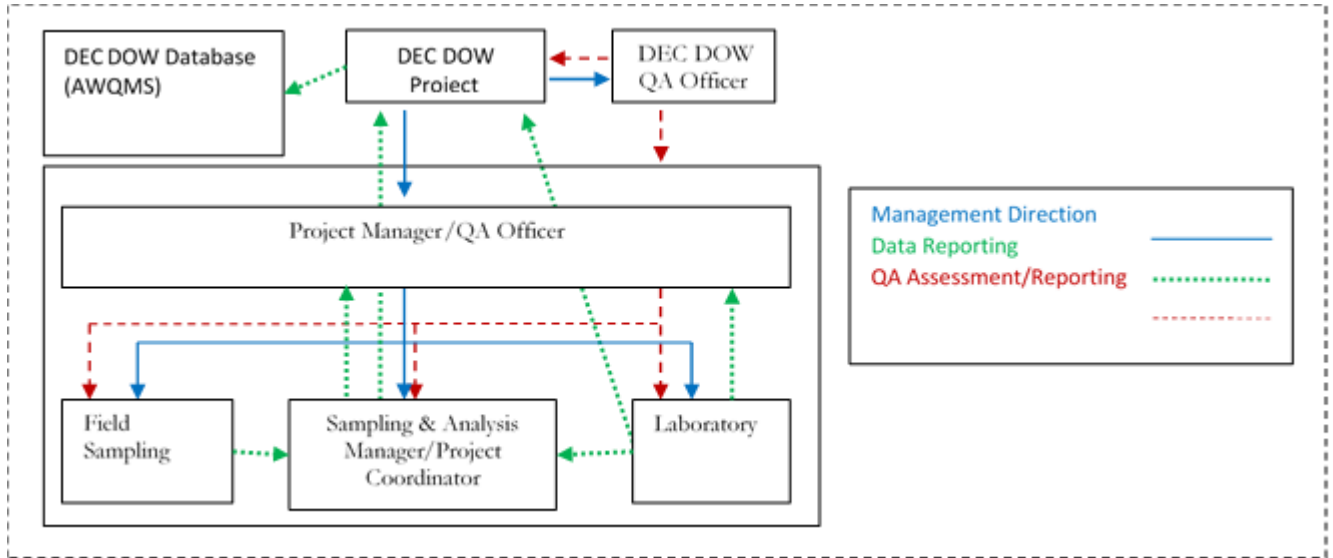


Figure 1. Project Organizational Structure

A.4 Project Definition/Background and Project Objectives

A.4.1 Project Definition

DEC identified the following beaches through a public nomination process; the Recreational Beach Survey is available at: <http://dec.alaska.gov/water/wqsar/wqs/beachprogram.htm>. The qualifying Hoonah Beaches are listed below.¹

- Inner Point Sophia (58.12952, -135.464858)
- Gartina Harbor Way (Gus's Beach 58.105764, -135.443018; Shellfish harvest area 58.104790, -135.450697; Harbor 58.104976, -135.441066)

Based on the information provided by respondents, DEC ranked these beaches as Tier 1. Tier 1 includes high priority beaches that pose the greatest threat of human contact with contaminated waters during recreational use. Contact with waters containing fecal contamination increases the risk of becoming ill due to pathogens contained in feces.

¹ One replicate of each analysis will be taken each at sampling event.

A.4.2 Project Background

The Beaches Environmental Assessment and Coastal Health (BEACH) Act was passed by the U.S. Congress in 2002 in response to increased occurrences of water-borne illnesses at recreational beaches. The EPA administers grant funds to states, tribes, and territories under the BEACH Act to establish monitoring and public notification programs, such as the Alaska BEACH Program. The BEACH program has established national marine water quality monitoring and reporting standards for fecal waste contamination and notifies the public when levels exceed state standards. The Hoonah beaches are some of multiple locations monitored under the Alaska BEACH Program².

A.4.3 Project Objective(s)

The objectives for this project are to:

- Monitor selected beaches for fecal indicator organisms (i.e., fecal coliform and enterococci bacteria) during periods of high recreational use.
- Notify the public when indicator organisms exceed Alaska Water Quality Standards (WQS).

The first objective will be achieved through designing a monitoring plan that samples at identified beaches during periods of high recreation activity by the public³.

The secondary objective will be achieved by distributing data to the public and stakeholders through DEC's Beach Program webpage, posters, press release and social media posts.

A.5 Project/Task Description and Schedule

A.5.1 Project Description

The HIA Project Liaison, HIA Lead Field Sampler will collect water samples from the Hoonah beaches during the 2022 recreation season. A DEC-approved laboratory will analyze samples for presence of fecal coliforms by SM 9222 D and Enterococci by ASTM D6503. Monitoring results will be posted on the DEC Beach Program webpage and distributed to stakeholders through a listserv.

A.5.2 Project Implementation Schedule

Table 3 includes the implementation schedule and sampling frequency for selected parameters and methods.

² Beaches being monitored in 2022 as part of the Alaska BEACH Program include Ketchikan, Hoonah, and Kenai River beaches.

³ High use periods for this project are defined as ice-free months between May and September, with the highest use periods occurring in June through August.

Table 2: Implementation Schedule for Selected Parameters and Methods

Product	Measurement/ Parameter(s)	Sampling Site	Sampling Frequency	Time Frame
Field Sampling	Ambient air temperature, turbidity (Hach Turbidimeter), pH, conductivity, marine water temperature (Hanna handheld meter)	All sites	Each sample event	June - September
Lab Analysis	Fecal coliform (SM 9222 D), Enterococci (ASTM D6503), MST Human, Dog, Gull Bacteroides	All sites	Each sample event	June - September
Field Audit	Audit of field monitoring operations	All sites	At least once per field season	June - September
Field Replicate	Fecal coliform (SM 9222 D), Enterococci (ASTM D6503)	One site, alternate	Each sample event	June - September

A.6 Data Quality Objectives and criteria for Measurement Data

A.6.1 Data Quality Objectives (DQOs)

Data Quality Objectives (DQOs) are qualitative and quantitative statements which are derived from the DQO Process that:

- Determine ambient water concentrations of pathogens (fecal coliforms and enterococci) and compare these values to marine water quality standards (18 AAC 70).
- The goal of the project is to monitor two Tier 1 beaches to determine current conditions and if the beaches meet regulatory limits for contact recreational marine water quality standards. The measurement system is designed to produce water pollutant concentration data that are of the appropriate quantity and quality to assess compliance with state water quality standards at a screening level assessment.

A.6.2 Measurement Quality Objectives (MQOs)

Measurement Quality Objectives (MQOs) are a subset of DQOs. MQOs are designed to evaluate and control various phases (sampling, preparation, and analysis) of the measurement process to ensure that total measurement uncertainty is within the range prescribed by the project’s DQOs. MQOs define the acceptable quality (data validity) of field and laboratory data for the project. MQOs are defined in terms of the following data quality indicators:

- Detectability
- Precision
- Bias/Accuracy

- Completeness
- Representativeness
- Comparability

Detectability is the ability of the method to reliably measure a pollutant concentration above background. DEC DOW uses two components to define detectability: method detection limit (MDL) and practical quantification limit (PQL) or reporting limit (RL).

- The MDL is the minimum value which the instrument can discern above background but no certainty to the accuracy of the measured value. For field measurements the manufacturer’s listed instrument detection limit (IDL) can be used.
- The PQL or RL is the minimum value that can be reported with confidence (usually some multiple of the MDL).

Note: The measurement method of choice should at a minimum have a practical quantification limit or reporting limit 3 times more sensitive than the respective DEC WQS and/or permitted pollutant level (for permitted facilities).

Sample data measured below the MDL is reported as ND or non-detect. Sample data measured \geq MDL but \leq PQL or RL is reported as estimated data. Sample data measured above the PQL, or RL is reported as reliable data unless otherwise qualified per the specific sample analysis.

Precision is the degree of agreement among repeated measurements of the same parameter and provides information about the consistency of methods. Precision is expressed in terms of the relative percent difference (RPD) between two measurements (A and B).

For field measurements, precision is assessed by measuring replicate (paired) samples at the same locations and as soon as possible to limit temporal variance in sample results. Field and laboratory precision is measured by collecting blind (to the laboratory) field replicate or duplicate lab samples. For paired and small data sets project precision is calculated using the following formula:

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

For larger sets of paired precision data sets (e.g., overall project precision) or multiple replicate precision data, the following formula may be used:

$$RSD = 100 * (\text{standard deviation} / \text{mean})$$

Note: Precision assessed only when both paired values \geq :

- 5 times PQL (fecal coliforms SM 9222D)
- 2 times PQL (E. coli SM 9222B)

Bias (Accuracy) is a measure of confidence that describes how close a measurement is to its “true” value. Methods to determine and assess accuracy of field and laboratory measurements include, instrument calibrations, various types of QC checks (e.g., sample split measurements, sample spike recoveries, matrix spike duplicates, continuing calibration verification checks, internal standards, sample

blank measurements (field and lab blanks), external standards), performance audit samples (DMRQA, blind Water Supply or Water Pollution PE samples from A2LA certified, etc.), Bias/Accuracy is usually assessed using the following formula:

$$Accuracy = \frac{Measured\ Value}{True\ Value} \times 100$$

Completeness is a measure of the percentage of valid samples collected and analyzed to yield sufficient information to make informed decisions with statistical confidence. As with representativeness, data completeness is determined during project development and specified in the QAPP. Project completeness is determined for each pollutant parameter using the following formula:

$$\frac{T - (I + NC)}{T} \times 100\% = Completeness$$

- Where: T = Total number of expected sample measurements.
I = Number of invalid samples measured results.
NC = Number of sample measurements not produced (e.g., spilled sample, etc.).

This project has a goal of 80% data completeness. In 2022, 13 sampling events are planned. The 2022 monitoring data is intended for a screening level assessment only in future IR cycles. The data collected is intended to provide members of the public with pertinent recreation information.

Representativeness is determined during project development and specified in the QAPP. Representativeness assigns what parameters to sample for, where to sample, type of sample (grab, continuous, composite, etc.) and frequency of sample collection.

Comparability is a measure that shows how data can be compared to other data collected by using standardized methods of sampling and analysis.

Monitoring shall be conducted in accordance with EPA-approved analytical procedures by state certified or equivalent laboratories and in compliance with 40 CFR Part 136, Guidelines Establishing Test Procedures for Analysis of Pollutants, as listed in Table 4. Field parameters will be measured using a Hach turbidimeter, HANNA® handheld probe, or an equivalent sonde (minimum resolution of 0.1 °C or better) as a point measurement. The device used must be verified prior to each sampling event⁴.

Each sampling location is fixed and located by a GPS coordinate. The locations do not change throughout the sampling season, but the area of sampling may change due to targeted parameters during a sampling event. Sampling is conducted in accordance with the Hoonah BEACH Monitoring Handbook (ADEC 2022.a).

⁴ See Appendix B: Standard Operating Procedure for Ambient Water Collection for Pathogen Monitoring and the Hoonah BEACH monitoring handbook for more information on equipment calibration and maintenance schedules.

Table 3: Project Measurement Quality Objectives

Group	Analyte	Method	MDL	PQL	Precision (RPD)	Accuracy
Pathogens	Fecal coliform	SM 9222 D, Membrane filtration (MF)	1 CFU/100 mL	1 CFU/100 mL	±60%	NA
	Enterococci	D6503-99, Enterococci by Enterolert	1 MPN/100 mL	1 MPN/100 mL	±60%	NA
	Microbial Source Tracking	Polymerase Chain Reaction	NA	NA	NA	NA
Field	Temperature Air/Water	EPA 170.1	NA	0.1°C	±0.02°C	± 0.2 °C
	pH	EPA 150.2	NA	NA	NA	±0.05 pH
	Turbidity	EPA 180.1, ASTM D1889, SM 2130 B	NA	NA	± 1% of reading or 0.01 NTU, whichever is greater	± 1 NTU

A.6.3 Data Validation and Verification

All data generated shall be validated in accordance with the QA/QC requirements specified in the methods and the technical specification outlined in this QAPP. Raw field data will be maintained by the Program staff who collect it. Raw laboratory data shall be maintained by the laboratory. The laboratory may archive the analytical data into their laboratory data management system. All data will be kept a minimum of 5 years.

The summary of all laboratories analytical results will be reported to the Project Coordinator/Manager staff. Protocols for laboratory data validation and verification are listed in Section B.4.2 and as specified in the laboratory's QAPP and SOPs.

Staff will verify that equipment used to collect field data is reading within acceptable limits before each sampling event using calibration solution. After sampling is completed, staff will complete a post verification check on equipment using calibration solution. Staff will record the date, name of equipment operator, calibration solution lot number and expiration date, reading of the standard solution, and verification pass/fail in a logbook kept with the field instrument.

Unacceptable data (i.e., data that do not meet the QA measurement criteria of precision, accuracy, representativeness, comparability, and completeness) will not be used for further analyses but will be documented. Any problems with the data will be clearly defined, flagged appropriately and data use clearly delimited and justified. Any action taken to correct QA/QC problems in sampling, sample handling, and analysis must be noted. Under the direction of the Project Manager/Coordinator, project staff **will document all QA/QC corrective actions taken.**

The Project Coordinator or his/her designee is responsible for reviewing electronic or paper data sheets for accuracy and completeness within 48 hours of each sample collection activity, if possible. The Project Coordinator or his/her designee will compare the sample information in the electronic or paper field sheets with the laboratory analytical results to ensure that no transcription errors have occurred, and to verify project QC criteria have been met (e.g., samples preserved, and sample hold times met as required by QAPP and method, relative percent difference (RPD) results for blind sample replicates).

RPD's greater than the project requirements will be noted. The Project Manager/Coordinator, along with supervisors and/or the Project QA Officer, if necessary, will decide if any QA/QC corrective action will be taken if the precision, accuracy (bias) and data completeness values exceed the project's MQO goals.

The Project Manager/Coordinator and the QA/QC Officer will review and validate data against the Project's defined MQOs prior to final reporting stages. If there are any problems with quality sampling and analysis, these issues will be addressed immediately, and methods will be modified to ensure that data quality objectives are being met. Modifications to monitoring will require notification to DEC and subsequent edits to the approved QAPP.

Only data that have been validated and qualified, as necessary, shall be provided to DEC Division of Water and entered in the applicable database (AWQMS, WQX).

A.7 Special Training Requirements/Certification

NPS Section Manager is responsible for overall technical and contractual management of the project. The current NPS Section Manager is up to date on current management training(s) and has over 20+ years' experience in the NPS Section.

DEC Hoonah beaches Project Manager is responsible for overall technical and contractual management of the project. She currently serves as DEC DOW's BEACH Grant coordinator and has experience in administrating BEACH Grant Monitoring Program grants. The experience associated with their duties allows them to be effective in carrying out duties as Project Manager.

SAWC Project Manager is responsible for coordinating efforts for field sampling, including equipment and supplies procurement, planning and leading field sampling events. The Project Manager is also responsible for preliminary QAQC of field data.

HIA Project Lead Field Sampler will assist the SAWC Project Manager with field sampling, including equipment and supplies procurement, equipment maintenance, data organization, and other tasks as needed.

Project QA Officer is the DEC DOW's Quality Assurance Officer. His training and experience allow him to successfully fulfill his duties as Project QA Officer.

Subcontracted Laboratory performing analytical work must have the requisite knowledge and skills in execution of the analytical methods being required. Information on laboratory staff competence is usually provided in each lab’s Quality Management (QMP) and/or Quality Assurance Plan (QAP). The laboratories to be used during the 2022 field season, will be Alaska Drinking Water certified microbiological laboratories, or maintain equivalent certification. It is the responsibility of the contracted lab to maintain a current copy of the laboratory’s QA Plan and attendant method specific SOPs on file with the NPS Section Manager, Project Manager, and DEC DOW QA Officer during the duration of laboratory use.

NPS Section Manager: Laura Eldred

Project Manager: Gretchen Augat

DEC QA Officer: John Clark

SAWC Project Manager/QA Officer: Rebecca Bellmore, Science Director; Rob Cadmus, Executive Director

HIA Project Lead Field Sampler: Jesse Endert, Environmental Specialist

HIA Field Support Personnel: TBA

<i>Table 4: Training Requirements</i>				
Specialized Training/Certification	Field Staff	Project Manager	NPS Section Manager	Project QA Officer
Safety training	X	X	X	X
Water sampling techniques	X	X	X	X
Instrument calibration and QC activities for field measurements	X	X		X
QA principles				X
Chain of Custody procedures for samples and data	X	X		X

A.8 Documents and Records

Paper field data sheets will be provided to all field crews (Appendix A). An electronic tablet may be used to digitally record field measurement as well⁵. The lead field sampler is responsible for ensuring that all field data forms are correct.

Field activities and observations will be recorded on paper data sheets (or electronic, if available). Any comments or descriptions will be noted in the comments with enough detail so that participants can reconstruct events later if necessary. Survey results and field data sheets will include descriptions of any changes at the site personnel and responsibilities or deviations from the QAPP/SAP as well as the

⁵ In 2021 the EPA released a digital version of the BEACH Sanitary Survey. The digital version is accessed through ESRI 123 App. The survey may be used for the 2022 monitoring season; however, paper field sheets should still be filled out as backup documentation.

reasons for the changes. Requirements for the field survey and field data sheet entries will include the following:

- Entries will be made while activities are in progress or as soon afterward as possible (the date and time that the notation is made should be included, as well as the time of the observation itself).
- Each entry will have its own unique identifier for the sampling event.
- Unbiased, accurate language will be used.
- If paper copies of the field data sheets are submitted, entries will be made legibly with **black (or dark) waterproof ink**.
 - Data or other information that has been entered incorrectly will be corrected by drawing a line through the incorrect entry and initialing and dating the lined- through entry. Under no circumstances should the incorrect material be erased, made illegible or obscured so that it cannot be read.
- Any deviation from the sampling plan will be included in the comments of the field data form.
- When field activity is complete, the electronic field survey form will be submitted and saved to the digital project file.

In addition to the preceding requirements, the person recording the information must have an additional field crew member review the data entry, either on the electronic survey application or the paper copy, of the data sheets. After data review is complete, the field crews will submit the data electronically to the database (e.g., AWQMS, WQX, EPA BEACON). The Project Manager will conduct the first round of quality assurance reviews, including field and laboratory datasets. Staff will then request a QA review from the NPS Section Manager. The data will then be submitted to the QA Officer for review. After the final QA review is completed, data will be uploaded electronically into AWQMS. The type of information that may be included in the electronic survey and/or paper field data forms includes the following:

- Names of all field staff
- A record of site health and safety meetings, updates, and related monitoring
- Station name and location
- Date and collection time of each sample
- Observations made during sample collection, including weather conditions, environmental conditions, complications, potential bacteria sources, and other details associated with the sampling effort
- Photo log⁶

Field logbooks/field data sheets and sample chain-of-custody forms will be completed for all samples and kept in the project file. Laboratory data results from the laboratories are recorded on laboratory data sheets, bench sheets and/or in laboratory logbooks for each sampling event. These records as well

⁶ See **Error! Reference source not found.** for specific information on data requirements for drone use.

as control charts, logbook records of equipment maintenance records, calibration, and quality control checks, such as preparation and use of standard solutions, inventory of supplies and consumables, check in of equipment, equipment parts and chemicals are kept on file at the laboratory.

Any procedural or equipment problems are recorded in the field notebooks/field data sheets. Any deviation from this Quality Assurance Project Plan will also be noted in the field notebooks/field data sheets. Data results will include information on field and/or laboratory QA/QC problems and corrective actions.

In addition to any written report, data collected for the project will be provided electronically in an AQWMS compatible format, which will be provide by DEC.

All records will be retained according to state records retention schedule. Table 6 includes a description of types of records/document types that may be included.

<i>Table 5: Project Documents and Records</i>	
Categories	Record/Document Types
Site Information	Site maps
	Site pictures
Environmental Data Operations	QA Project Plan
	Field Method SOPs
	Field Notebooks/Field Data Sheets
	Sample collection/measurement records
	Sample Handling & Custody Records
Raw Data	Inspection/Maintenance Records
	Lab data (sample, QC, and calibration) including data entry forms
Data Reporting	Progress reports
	Project data/summary reports
	Lab analysis reports
Data Management	Data quality assessments
	Site audits
	Lab audits
	QA reports/corrective action reports
	Corrective Action Response

B. Data Generation and Acquisition

B.1 Sampling Process Design (Experimental Design)

Monitoring will be conducted at preselected locations.

In 2022, monitoring will occur 13 times throughout the open water season (approximately June through early September) to capture conditions at different hydrological regimes and human use levels.

Sampling will occur twice in June, July, and August, and once in early September. The sampling schedule is available upon request.

Water samples will be analyzed to determine the population densities of microbes that indicate the presence of fecal contamination; microbes to be enumerated will be enterococci and fecal coliforms, with the results reported per 100 mL marine water.

For each sample collected, the date and time will be noted. Sample containers will be delivered to labs for analysis within the six (6)⁷ hour hold time required for pathogens for accurate results.

B.1.1 Define Monitoring Objective(s) and Appropriate Data Quality Objective(s)

Project schedule and tasks may be adjusted as needed due to unplanned or unavoidable events.

MONITORING PROJECT TASKS

TASK 1: Review 2020 Hoonah Beach Sampling Plan and Quality Assurance Project Plan (SAP/QAPP) and Beach Monitoring Handbook

Deliverable(s) and Permits: Review and comment on the DEC-provided Beach Sampling Plan and Quality Assurance Project Plan (SAP/QAPP) and Beach Monitoring Handbook. SAWC will work with the DEC project manager to update the 2021 Hoonah Beach SAP/QAPP and Beach Monitoring Handbook and have signed by the relevant parties.

Deliverable	Due Date:
Comments/revisions on DEC-provided QAPP and Handbook (Word track changes)	March 31, 2022
Final QAPP and Beach Monitoring Handbook signed by all parties	May 1, 2022

TASK 2: Monitor Beach Water Quality

Deliverable(s) and Permits: SAWC will conduct marine water quality monitoring 13 times during the 2022 recreational use season (June to early September) for bacteria at recreational beaches using the DEC-approved SAP/QAPP.

During each sampling event SAWC will collect one (1) marine water sample at each beach location for fecal coliform bacteria (SM 9222D) and enterococci (ASTM D6503-99) using the DEC- approved SAP/QAPP and submit to a DEC-approved laboratory within the six (6) hour holding time. Collect one (1) replicate sample for each bacteria analytical test per sampling event for quality assurance. Through previous beach monitoring efforts, the microbial source has been identified and therefore additional MST sampling is not needed for the Hoonah beach monitoring locations during this recreational season.

The bacteria samples will be collected and submitted to the DEC-approved Admiralty Environmental, Inc. laboratory in Juneau for analytical testing. SAWC may collect the samples directly or contract with the Hoonah Indian Association, City of Hoonah, or another qualified entity to collect samples. Samples will be taken from the locations outlined in the SAP/QAPP. The 2 beaches monitored will include Inner Point Sophia and Gartina Harbor Way. One (1) QA laboratory sample per analyte will be collected per

⁷ Max hold time for pathogen samples is: 6 hours in the field, 2 hours in the laboratory (total of 8 hours hold time).

sampling event (alternating between sampling locations). SAWC will work with property owner(s) to secure permission to access land for sampling purposes. Field technicians will complete the project-specific field datasheet, and field parameters collected by a DEC-provided portable meter in the field at each sampling location during each sampling event, chain-of-custody forms, and site photos at each monitoring location for each monitoring event. Field technicians will take photos at each site during each sampling round to capture beach conditions.

Admiralty lab will analyze marine water samples collected by the field technicians from the 2 monitoring sites for fecal coliform bacteria (SM 9222D) and enterococci (ASTM D6503-99). One (1) replicate sample for each analytical test per sampling event will be analyzed for quality assurance. Approximately 39 samples will be analyzed for fecal coliform bacteria and enterococci. Admiralty lab will submit analytical results via email directly to DEC, SAWC and sample collection group. If a confirmed exceedance occurs, DEC will issue a beach advisory (formats include press release, DEC website, social media, list serv, radio and/or newspaper), and SAWC will conduct notification outreach, as necessary.

Sampling will occur twice in June, July, and August, and once in early September. The sampling schedule is available upon request.

Deliverable	Due Date:
Sanitary surveys from field datasheets (Excel and csv)	Within 36 hours of sampling event from June-September
Site photos (appropriate digital format)	
Chain of Custody form copies (PDF)	
Copy of other data information (e.g., calibration records)	
Land access permission documentation (if necessary)	

TASK 3: Educational Outreach

Deliverable(s) and Permits: Conduct educational outreach event to communicate the beach program findings following the recreational season. The typical prior recreational season outreach will be fulfilled in February 2022 (ACWA-B10-A3) during the 2021 monitoring program outreach which will include a discussion of 2022 recreational season plans and next steps for the program.

At the completion of sampling, SAWC will develop outreach material to communicate the beach program and sampling results to the Hoonah Community. Outreach material will be approved by DEC. SAWC will conduct outreach communication to the Hoonah community. Outreach will include public service announcements via radio, local newspaper and/or social media, and one area-appropriate presentation to the community.

Deliverable	Due Date:
Post-sampling outreach materials developed (electronic copies of draft presentations, press releases, etc.)	October 2022
Post-sampling outreach materials approved by DEC and communicated to the public (electronic copies of released material, list of attendees at presentations)	November 2022

TASK 4: Project Data Submission

Deliverable(s) and Permits: SAWC will compile and enter all monitoring data (e.g., analytical results, field parameters) into DEC-provided template which DEC will submit to three databases: Ambient Water Quality Monitoring System (AWQMS), Water Quality Portal (WQP), and Beach Advisory and Closing On-line Notification (Beacon). SAWC will create and submit a GIS geodatabase and map showing the spatial relationship between residential/public waste treatment and septic, topographic contours, surface water hydrology, potential pollution sources, and beach survey data; and provide the data in NAD83/Alaska Albers.

Deliverable	Due Date:
Monitoring data and results in DEC-provided electronic template (Excel workbook)	October 2022
GIS geodatabase and map	December 2022

TASK 5: Project Report

Deliverable(s) and Permits: SAWC will evaluate all sample results and submit a draft and final report of findings and conclusions. Report design must follow 2021 Hoonah Beach monitoring report prepared by SAWC for the 2021 recreational season.

The report will include background information, and the project need, objectives, and approach taken to meet the project objectives. The report will evaluate and describe project accomplishments, the environmental benefit and suggest future actions. Water quality analysis will use the DEC’s Listing Methodology for Determining Water Quality Impairments from Pathogens guidance, to compare results to the Marine Water Quality Indicator Criteria for bacteria. The report will include narrative description and tabular/graphical formats to evaluate the monitoring results. The report will include a quality assurance review describing the integrity of the reported analytical results as presented in the QAPP and data quality objectives. Appendices will incorporate all project data or refer to a web link and appropriate references.

Deliverable	Due Date:
Analyze results of beach monitoring	December 2022
Draft Beach Monitoring Report (Word)	January 2023
Final Beach Monitoring Report (Word and PDF)	February 2023

B.1.2 Identify the Site-Specific Sample Collection Location(s), Parameters to be Measured, and Frequencies of Collection

<i>Table 6: Site Location and Rationale</i>			
Site ID	Latitude	Longitude	Site Description
HB-InnerPtSoph	58.12952	-135.464858	Inner Point Sophia
HB-GartinaHbrWay	58.105222	-135.440927	Gartina Harbor Way Beach
Note: GIS maps of sampling locations (large scale as well as site specific) are shown in Appendix A.			

B.2 Sampling Method Requirements

Specific sampling methods are detailed in the Sampling SOP, included in the Appendix of this QAPP.

Laboratory samples will be listed as “grab” on the Chain-of-Custody forms and data sheets while field samples will be listed as “In situ” as defined below.

Grab Samples – Sample bottles will be filled sequentially, normally being filled to the shoulder of the bottle, leave a small space for expansion and mixing. The laboratory will provide sampling instructions with the sample bottles for specific samples.

In Situ Samples – In situ water measurements will be taken as point readings HANNA® Handheld probe. In situ measurements include air temperature and water temperature.

Turbidity – The sample bottles provided with the HACH® Turbidimeter will be used. Bottles will be rinsed with ambient water, and then filled to the level recommended by the manufacturer. Follow manufacturer’s instructions for operating the Turbidimeter. Check that all calibration standards are not expired before use.

B.2.1 Sample Containers and Equipment

The sample container, preservation, and holding time requirements are tabulated below:

<i>Table 7: Preservation and Holding Times for the Analysis of Samples</i>					
Analyte	Matrix	Container	Necessary Volume	Preservation and Filtration	Maximum Holding Time
Temperature Air and Water, pH	Surface Water	N/A, direct measurement			
Turbidity	Surface Water	G ⁸	15 mL	Collect measurement in field, do not freeze vial	N/A. Collect measurement at field site
Fecal coliform	Surface Water	PA	150 mL	Cool < 4°C, do not freeze, Na ₂ SO ₃ preserved	8 hours total, (6 hrs. field, 2 hrs. lab)
Enterococci	Surface Water	PA	150 mL	Cool < 4°C, do not freeze, Na ₂ SO ₃ preserved	8 hours total, (6 hrs. field, 2 hrs. lab)
MST	Surface Water	PC	500 ml	Cool <10°C; do not freeze, unpreserved	48 hours
Notes: G = glass, PA = autoclavable plastic, PC = polycarbonate					

B.3 Sample Handling and Custody Requirements

B.3.1 Sample Custody Procedures

Samples and sample containers will be maintained in a secure environment from the time the bottles leave the field until the samples are received at the laboratory. The laboratory will maintain custody of bottles and samples using their normal custody procedures.

Samples must be in the sampler's possession or in a cooler sealed with signed and dated friable evidence tape on opposing sides of the cooler. When the cooler is sealed, the method of securing the samples must be such that tampering with samples or bottles is not possible. The cooler must be secured so that the lid cannot be removed without breaking the evidence tape or cutting the lock.

Transfer of samples will be accomplished using the laboratory's Chain-of-Custody (COC) form. When samples are transferred between personnel, such transfer will be indicated on the COC form with signature, date, and time of transfer. The COC will remain with the samples, sealed inside the cooler, until received by the laboratory. DEC will provide a copy of the contracted lab COC for staff to use during field work.

⁸ A set of 15 mL glass vials are provided with the HACH® Turbidimeter kit. Check that vials are not scratched, broken or missing before the field season.

If custody is broken at any time during sample transfer, a note must be made on the COC form accompanying the sample. Upon receipt at the laboratory, the laboratory sample custodian will make note if a breach of custody has occurred (for example, if a custody seal has broken during transport).

B.3.2 Shipping Requirements

Packaging, marking, labeling, and shipping of samples will comply with all regulations promulgated by the U. S. Department of Transportation in 49 CFR 171-177. Staff should receive the necessary training for shipping samples or consult with the contracted laboratory for shipping instructions.

Samples collected in plastic bottles may be placed in the cooler with sufficient padding (e.g., bubble wrap, cardboard, etc.) to limit movement of the bottles in the cooler during transport. The sealed plastic bags and plastic sample bottles will be placed into a cooler with gel-ice/blue-ice in plastic bags to maintain a temperature of <4 °C. A temperature blank, 250 or 500 mL in size, will be placed in the cooler. Temperature will be measured prior to shipment and upon receipt at the lab. The chain of custody (COC) form will be placed in a plastic bag within the cooler. The cooler will be taped closed securely using packing tape at the last sampling site. If the cooler is being transported by the field crew member directly to the laboratory, tape is not mandatory.

<i>Table 8: Sample Transport Chain Information</i>						
Business Type	Name	Address	Hours	Contact Information	Transport Leg	Estimated Transit Time
Deliver directly to flight	Alaska Seaplanes	Hoonah airport	Flight 202 – 8:00 am daily; Flight 204 11:00 am daily	(907) 789-3331	Daily flight	20 minutes
Lab couriers deliver to laboratory	Admiralty Environmental lab	641 W. Willoughby Ave. Suite 301 Juneau, AK 99801	8:00 am – 5:00 pm; last sample drop off without additional charge at 3:30 pm	(907) 463-4415	Motor Vehicle	30 minutes
Ship FedEx	Source Molecular Corporation (MST samples only)	4985 SW 74 th Court, Miami, FL 33155		(786) 220-0379	FedEx First overnight	48 hours

B.4 Analytical Methods and Requirements

Water quality analytical methods that will be used throughout this project are outlined below. All analysis methods used for this program are EPA-approved. The contracted laboratory will be a currently

DEC Drinking Water -certified laboratory, though the lab will be using methods specified for water/wastewater analysis. The contracted laboratory's current Quality Assurance Plan will be on file with DEC Division of Water Quality Assurance Office detailing their quality assurance procedures. Laboratory turnaround time is 36 hours. Any issues regarding analytical data quality will be resolved by the DEC project manager in consultation with any or all the following: DEC QA Officer, sampling staff and the laboratory project manager.

B.4.1 Sampling Parameters

- **Temperature** will be reported in °C for air and water and will be measured using a Hanna handheld meter or an equivalent meter (minimum resolution of 0.1 °C or better). The thermometer will have current NIST traceable certification.
- **pH** will be collected using a HANNA handheld meter or similar device. Equipment must be calibrated prior to sampling season.
- Turbidity will be measured using a portable turbidimeter provided by ADEC. Fill provided vial following the manufactures instructions. Device must be calibrated prior to sampling event.
- **Fecal Coliform** Standard Method 9222D will be used to determine the fecal coliform concentration in surface water. Filter sample through a membrane filter. Place membrane on mFC agar containing aniline blue as indicator. Incubate at 44.5°C for 22-24 h. Colonies that are various shades of blue are positive for fecal coliforms. The blue color indicates the capability to ferment lactose to acid.
- **Enterococci** ASTM Method D6503-99 will be used to determine the most probable number enterococci concentration in surface water. Add reagent to the sample, pour into Quanti-Tray® or Quanti-Tray® /2000, seal in Quanti-Tray® Sealer and incubated for 24 hours at 41°C. Count fluorescent wells and refer to most probable number table.
- **Fecal Coliform** Standard Method 9221A SM9221 E (2) with A-1 media, MPN, marine growing waters method. This method describes multiple-tube fermentation procedures [also called the most probable number (MPN) procedure] for the detection and enumeration of fecal coliform bacteria in biosolids. These methods use culture-specific media and elevated temperature to isolate and enumerate fecal coliform organisms.

Monitoring shall be conducted in accordance with EPA-approved analytical procedures and in compliance with 40 CFR Part 136, Guidelines Establishing Test Procedures for Analysis of Pollutants. Reference the Project's MQO **Error! Reference source not found.**(section **Error! Reference source not found.**) of this QAPP for list of parameters of concern, approved analytical methods, method-specific detection and reporting limits, accuracy and precision values applicable to this project. 40 CFR, Part 136.6 lists other regulated pollutant parameters not listed in the MQO **Error! Reference source not found.**(section **Error! Reference source not found.**).

An expedited reporting turnaround time after sampling will be required for laboratory microbiological analyses to obtain results quickly for decision-making purposes. As pathogen exposure remains a risk to beach users during the period between sample analysis and reporting sample results, a short reporting time is recommended; a period of 36 hours following sample

submission should be used for reporting results to the QAO, the BPM, and local community point of contact.

Microbial Source Tracking Detection of the fecal associated Human, and other selected host species, gene biomarker by real-time quantitative Polymerase Chain Reaction (qPCR) DNA analytical technology. Each submitted water sample was filtered through 0.45-micron membrane filters. Each filter was placed in a separate, sterile 2ml disposable tube containing a unique mix of beads and lysis buffer. The sample was homogenized for 1 min and the DNA extracted using the Generite DNA-EZ ST1extraction kit (GeneRite, NJ), as per manufacturer's protocol.

Amplifications to detect the target gene biomarker were run on an Applied Biosystems StepOnePlus real-time thermal cycler (Applied Biosystems, Foster City, CA) in a final reaction volume of 20ul containing sample extract, forward primer, reverse primer, probe and an optimized buffer. The following thermal cycling parameters were used: 95°C for 10 min and 40 cycles of 95°C for 15 s and 60°C for 1 min. All assays were run in duplicate.

For quality control purposes, a positive control consisting of horse fecal DNA and a negative control consisting of PCR-grade water, were run alongside the sample(s) to ensure a properly functioning reaction and reveal any false negatives or false positives. The accumulation of PCR product is detected and graphed in an amplification plot. If the fecal indicator organism is absent in the sample, this accumulation is not detected, and the sample is considered negative. If accumulation of PCR product is detected, the sample is considered positive.

B.5 Quality Control Requirements

Table 4 lists the relative percent difference of field and laboratory replicates to be used for quality control (see section A.6.2 for discussion on calculation of precision and accuracy). The precision of field and laboratory measures will be calculated using the equation in section A.6.2. Data measurements that do not meet the limits described in A.6.2 may or may not be used in the final report depending on degree to which limits are not met. However, the report will clearly flag all data of questionable value along with a brief description of the problem and any justification why data should be considered for use. Daily field records (a combination of electronic and paper field data sheets) will make up the main documentation for field activities. As soon after collection as possible, field notes, data sheets, and chain-of-custody forms will be scanned to create an electronic record. Field data will be hand-entered into the database.

An example Data Management Flow Chart (Figure 2) provides a visual summary description of the data flow/management process for environmental data collected in support of DEC's Division of Water decision making processes. Revisions may be made as appropriate for the monitoring project.

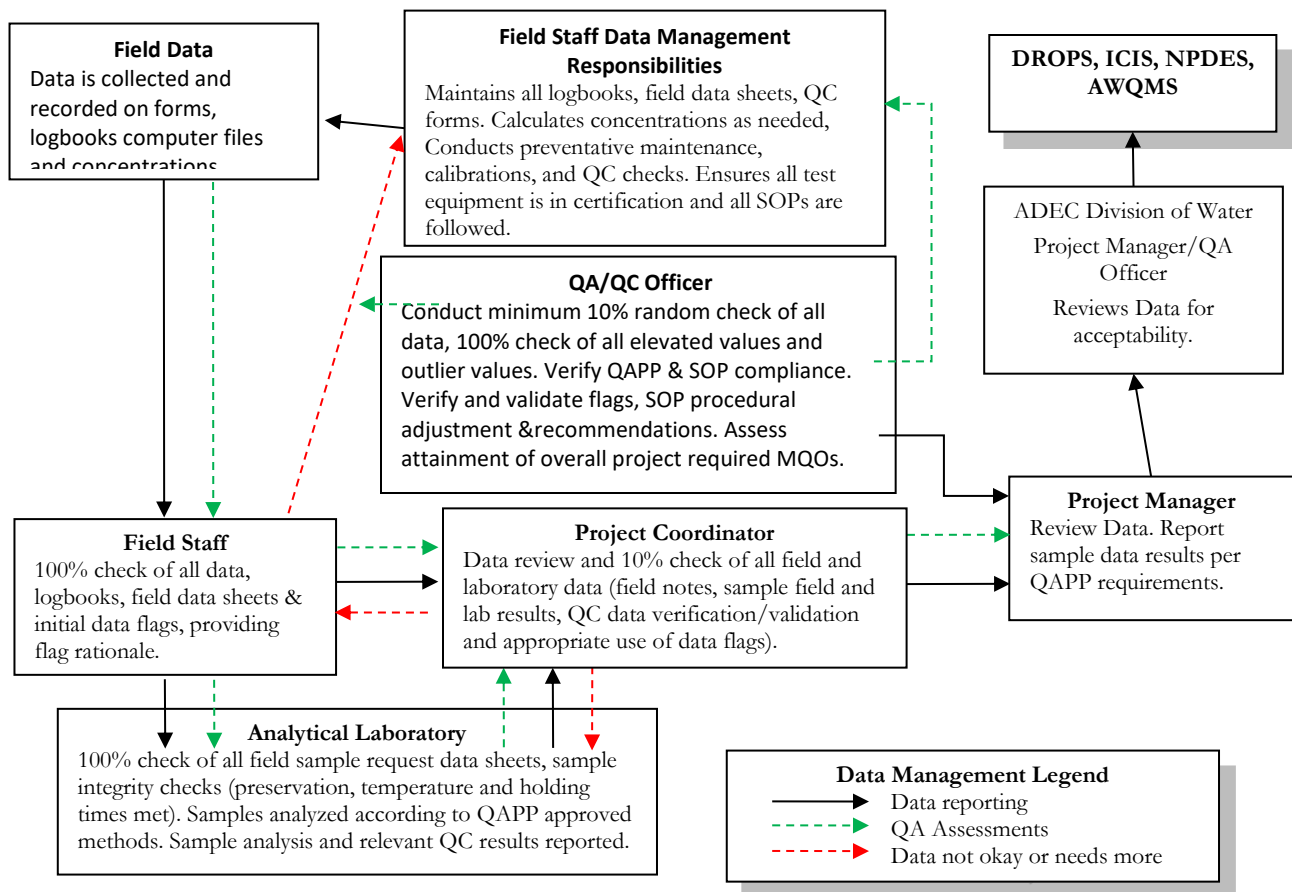


Figure 2. Data Management Flow Cart

One field sample replicate (i.e., duplicate) will be collected once each sampling event, at alternating sample locations, for both Fecal coliform and enterococci bacteria. The purpose of field sample replicate is to assess sampling and laboratory precision for the monitoring project.

For laboratory analyses, contract laboratories will submit quality control results along with sample analytical results. Laboratory Quality Control will include duplicates, holding times, sample temperatures upon receipt of sample at lab and blanks. Laboratory precision criteria should be within MQO criteria provided in Section A.6.

B.5.1 Field Quality Control

Quality control activities in the field will include adherence to documented procedures and the comprehensive documentation of sample collection information included in the field survey (electronic or paper). A rigidly enforced chain-of-custody program will ensure sample integrity and identification. The chain-of-custody procedure documents the handling of each sample from the time the sample was collected to the arrival of the sample at the laboratory.

Quality Control measures in the field include but are not limited to:

- Proper handling of sampling equipment.

- Maintenance, cleaning, and calibration of field equipment/ kits per the manufacturers and/or laboratory's specifications, and field Standard Operating Procedures (SOPs).
- Chemical reagents and standard reference materials are used prior to expiration dates.
- Proper field sample collection and analysis techniques, including but not limited to: Utilization of clean hands, dirty hands techniques; use of powder free nitrile gloves.
- Ensuring all sample equipment and sample containers are in proper condition (i.e., no cracks or broken bottle caps, tamperproof seals are intact before sampling)
- Correct sample labeling and data entry.
- Proper sample handling and shipping/transport techniques, including the use of a temperature blank in each cooler containing samples to be shipped.
- Field replicate measurements at a minimum of one sample for each analyte per sampling event.

Analytical methods used on the project have been approved and documented by EPA, Standard Methods, or ASTM. These methods will be used as project-specific protocols to document and guide analytical procedures. Adherence to these documented procedures will ensure that analytical results are properly obtained and reported.

B.5.2 Laboratory Quality Control (QC) Measures

Contracted and sub-contracted laboratories will follow the testing, inspection, maintenance, and quality control procedures required by EPA Clean Water Act approved methods and as stated in the respective laboratory's QAP and SOPs including the following:

Laboratories detail QC procedures used in their laboratory Quality Assurance Plan and method specific SOPs Quality Control in laboratories includes the following:

- Laboratory instrumentation calibrated with the analytical procedure.
- Laboratory instrumentation maintained in accordance with the instrument manufacturer's specifications, the laboratory's QAP and Standard Operating Procedures (SOPs).
- Specific QC activities prescribed in the project's QAPP.
- Laboratory data verification and validation prior to sending data results to DEC.

Contracted and sub-contracted laboratories will provide analytical results after verification and validation by the laboratory QA Officer. The laboratory must provide all relevant QC information with its summary of data results so that the project manager and project QA officer can perform field data verification and validation and review the laboratory reports. The project manager reviews these data to ensure that the required QC measurement criteria have been met. If a QC concern is identified in the review process, the Project Manager and Project QA Officer will seek additional information from the sub-contracted laboratory to resolve the issue and take appropriate corrective action/s.

B.5.3 QA Reports to Management

Following field and laboratory quality control measurements, quality analysis reports will be filed with the Project Coordinator and/or Project Manager. Table 10 details the report requirements for submittal to the Project Coordinator and/or Project Manager.

<i>Table 9: QA Reports to Management</i>					
QA Report Type	Contents	Presentation Method	Report Issued by	Reporting Frequency	
				As Required	Annual
On-site Field Inspection Audit Report	Description of audit results, audit methods and standards/ equipment used and any recommendations	Checklist or written text and tables, charts, graphs displaying results	Project QA Officer/ Coordinator/ Manager	✓	
Corrective Action Recommendation	Description of problem(s); recommended action(s) required; time frame for feedback on resolution of problem(s)	Written text/table	Project QA Officer/ Coordinator/ Manager	✓	
Response to Corrective Action Report	Description of problem(s), description/date corrective action(s) implemented and/or scheduled to be implemented	Written text/table	Project Manager/ Coordinator overseeing sampling and analysis	✓	
Data Quality Audit	Independent review and recalculation of sample collection/analysis (including calculations, etc.) to determine sample result. Summary of data audit results; findings; and any recommendations	Written text and charts, graphs displaying results	Project QA Officer	✓	
Quality Assurance Report to Management	Project executive summary: data completeness, precision, bias/ accuracy	Written text and charts, graphs displaying results	Project QA Officer	✓	✓

B.6 Instrument Calibration and Frequency

Field instruments shall be calibrated prior to using the instruments. The Project Manager will ensure that instruments are calibrated correctly, and appropriate documents recorded and retained. Sensors for field equipment (i.e., air and water temperature) will be replaced according to manufacturer's

recommendations. If abnormal readings occur, the manufacturer will be contacted for assistance or replacement of field equipment.

Contracted and sub-contracted laboratories will follow the calibration procedures found in its QAP and the laboratory's Standard Operating Procedures (SOPs). Specific calibration procedures for regulated pollutants will agree with the respective "EPA Approved" Clean Water Act Pollutant methods of analysis. Field and/or Laboratory calibration records will be made available to DEC upon request.

B.7 Inspection/Acceptance of Supplies and Consumables

Pre-cleaned sample containers will be obtained from the lab with the appropriate preservation method included. Coolers, gel ice, temperature blanks, and chain-of-custody forms will be provided by the contract laboratory prior to field mobilization. Qualified staff will check all field equipment and supplies to ensure that their technical specifications have been met before use. Any deviations during inspection procedures will be remedied by the project manager and recorded in the electronic or paper field data sheets. If re-sampling becomes necessary, replacements will be made.

No standards, solutions, buffers, or other chemical additives will be used if the expiration date has passed. It is the responsibility of the sampling manager or his/her designee to keep appropriate records, such as logbook entries or field data sheets, to verify the inspection/acceptance of supplies and consumables and restock these supplies and consumables when necessary.

Contracted and sub-contracted laboratories will follow procedures in their laboratory's QAP and SOPs for inspection/acceptance of supplies and consumables.

B.8 Data Acquisition Requirements (Non-Direct Measurements)

Topographic non-direct measurements (e.g., maps, charts) will be conducted using USGS derived materials. All geographical materials will be listed according to their source, year, and scale. GPS information will be documented by including collection device make and model number, geographic coordinate system, degree of accuracy (minimum of three satellite signals), and calibration information. GIS information will include GIS software program and model, source information, and geographic coordinate system.

B.9 Data Management

Various people are responsible for separate or discrete parts of the data management process:

The field samplers are responsible field measurements/sample collection and recording of data and subsequent shipment of samples to laboratories for analyses. They assemble data files, which includes raw data, calibration information and certificates, QC checks (routine checks), data flags, sampler comments and metadata where available. These files are assembled and forwarded for secondary data review by the sampling supervisor.

Laboratories are responsible to comply with the data quality objectives specified in the QAPP and as specified in the laboratory QAP and method specific SOPs. Validated sample laboratory data results are reported to the sampling coordinator/supervisor/project supervisor.

Secondary reviewers (sampling coordinator/supervisor/project supervisor) are responsible for the QC the review, verification and validation of field and laboratory data and data reformatting as appropriate for reporting to AWQMS and reporting validated data to the project manager.

The project QA officer is responsible for performing routine independent reviews of data to ensure the monitoring projects data quality objectives are being met. Findings and recommended corrective actions (as appropriate) are reported directly to project management.

The project manager is responsible for final data certification.

DEC DOW project manager and QA Officer AQS data entry staff conducts a final review (tertiary review) and submits the validated data to AWQMS.

Daily field records (a combination of field and core logbooks data sheets) will make up the main documentation for field activities. As soon after collection as possible, field notes, data sheets, core logs, and chain-of-custody forms will be scanned to create an electronic record. Field data will be hand-entered into the database. One hundred percent of the transferred data will be verified based on hard copy records. Electronic QA checks to identify anomalous values will also be conducted following entry.

Data obtained during sampling activities will be entered into field notebooks.

The following is a list of data information that will be kept and submitted to DEC:

- Field equipment and chemicals maintenance, cleaning, and calibration records
- Field notebooks
- Sample Data Sheets
- Photographs of sampling stations and events
- Chain-of-Custody forms
- Laboratory equipment maintenance, cleaning, and calibration records
- Laboratory bench sheets, control charts, and SOPs
- Records of QA/QC problems and corrective actions (field and/or laboratory)
- Laboratory data QC records
- Records of data review sheets
- Replicate, performance evaluation records and other QA/QC control records (field and laboratory)
- Data review, verification, and validation records

Sample Numbering

All samples will be assigned a unique identification code based on a sample designation scheme designed to suit the needs of the field personnel, data management, and data users. Sample identifiers will consist of two components separated by dashes. The first component is used to identify the area to which the sample originated, for example: HB = Hoonah Beach.

Laboratory Data

The contract laboratory will submit data in electronic format to DEC. Written documentation will be used to clarify how field replicates and laboratory duplicates and QA/QC samples were recorded in the data meta tables and to provide explanations of other issues that may arise. The data management task will include keeping accurate records of field and laboratory QA/QC samples so that project managers and technical staff who use the data will have appropriate documentation. Data management files will be stored on a secure computer or on a removable hard drive that can be secured. All records will be retained by the contract laboratory for five years.

Data Storage and Retention

Data management files will be stored on a secure computer or on a removable hard drive that can be secured. Laboratory Records will be retained by the contract laboratory for a minimum of five years. Project records will be retained by the lead organization conducting the monitoring operations for a minimum of five years, preferably longer. Site location and retention period for the stored data will be specified in each QAPP.

C. Assessment and Oversight

C.1 Assessment and Response Actions

Assessment audits are independent evaluations of the monitoring project that are performed by the Project's QA Officer or his/her designee. These audits may include (but are not limited to) any of the following: on-site field surveillance, on-site laboratory audits, performance evaluation samples, blind sample duplicates/ replicates (precision samples), field split samples, data quality audits, and/or data reviews. The number and types of assessments are dependent upon the monitoring project's intended data uses.

C.1.1 On-Site assessments to be performed

- One on-site field audit will be completed to evaluate sampling protocols and survey techniques. Audits will evaluate whether procedures used for sample collection, preservation, shipping and hold times, and sample receipt at lab follow QAPP requirements.

C.1.2 Project Data Assessments

- Audits of Monitoring Data for reproducibility of results from recalculation/reconstruction of field/lab data.
- Calculation of monitoring project's overall achieved precision, accuracy, and data completeness compared to QAPP defined precision, accuracy, and data completeness goals. Method specific precision, accuracy, and data completeness criteria are specified in the Project MQO Table 3 of section A.6.2.
- Complete the data review checklist. Describes whether project data quality objectives and measurement quality objectives were obtained, and corrective actions that were taken if any.

- Water Quality Field Report will be completed at the end of the project. Summarizes project methods and results and whether exceedances of Alaska’s Water Quality Standards were measured.

C.2 Revisions to QAPP

This QAPP will be reviewed and revised annually or earlier as needed. Minor revisions may be made without formal comment. Such minor revisions may include changes to identified project staff, QAPP distribution list, and or minor editorial changes.

Revisions to the QAPP that affect state monitoring Data Quality Objectives, Method Quality Objectives, method specific data validation “critical” criteria and/or inclusion of new monitoring methods must solicit input/ and pre-approval by DEC DOW QA Officer/DEC Project Manager or Coordinator before being implemented.

D. References

- ADEC (Alaska Department of Environmental Conservation). 2022. Beach Monitoring Handbook, Hoonah, 2022 Monitoring Season. Water Quality, Division of Water. Juneau, AK. V.5.
- ADEC (Alaska Department of Environmental Conservation). 2020. 18 AAC 70. Water Quality Standards. 73 pg.
- Bureau of Land Management. 2017. AIM National Aquatic Monitoring Framework: Field Protocol for Wadeable Lotic Systems. Tech Ref 1735-2. U.S. Department of the Interior, Bureau of Land Management, National Operations Center, Denver, CO.
- USEPA (U.S. Environmental Protection Agency). 2013. Great Lakes Beach Sanitary Survey User Manual. EPA-823-B-06-001. U.S. Environmental Protection Agency, Office of Water. Washington, DC
- USEPA (U.S. Environmental Protection Agency). 2014. National Beach Guidance and Required Performance Criteria for Grant, 2014 Edition. EPA-823-B-14-001. U.S. Environmental Protection Agency, Office of Water. Washington, DC.

A. Appendix A: Sample Site Locations

A.1 2022 Monitoring Locations and site descriptions

Site ID	Latitude	Longitude	Site description
Inner Point Sophia	58.0739	-135.2743	Widely used recreation beach, swimming area, shellfish harvesting area, and cruise ship destination 1 mile from city center.
Gartina Harbor Way	58.0618	-135.2627	Primary shellfish, crab, fish harvesting area adjacent to Hoonah City Harbor.

A.2 2022 Sampling Location Maps



Figure A.2. Map of sampling locations at Inner Point Sophia Beach and Gartina Harbor Way Beach. Three sampling locations at Gartina Harbor Way Beach are shown in the insert.

B. Appendix B: Standard Operating Procedure for Ambient Water Collection for Pathogen Monitoring

B.1 Standard Operating Procedures Alaska BEACH Program

Sampling for the Alaska BEACH Program involves wading into the water adjacent to a beach commonly used for recreation to collect water from below the surface into sample jars. The sample should be collected in the general recreational beach area, or near locations expected to be influenced by fecal contamination (e.g., adjacent to sewage lagoons, near small boat harbors, etc.). Field staff will have completed sampling after the following steps have been accomplished:

- Each sample jar is filled with water,
- Each sample jar is labeled,
- Each sample jar is placed in a cooler kept chilled with artificial ice,
- The Beach Sampling Data Sheet is filled out,
- A chain-of-custody form is filled out,
- The cooler is transported to the laboratory responsible for determining fecal coliform and enterococcus populations,
- A copy of the Beach Sampling Data Sheet is sent to the respective Project Manager, and a copy of the Beach sampling Data Sheet is sent to Project Coordinator.

Detailed directions for collecting good water samples, shipping the samples to the laboratory, and providing beach assessment information to DEC are given in the following subsections.

B.1.1 Sample Collection Method

A good water sample is collected by avoiding cross-contamination, which can happen when the sampler inadvertently contaminates the sample. To reduce the potential for cross-contamination the sampler must follow a standard sample-collection method. Step-by-step sample-collection instructions are provided below:

1. Request a sample kit from the laboratory. The kit should include:
 - A cooler
 - The appropriate sample containers for marine water quality sampling (enterococcus and fecal coliform bacteria)
 - Artificial ice to keep the cooler chilled to the appropriate temperature
 - The appropriate container for the field blank
 - Temperature blank
 - Chain-of custody form

- Custody seals
 - Sample jar labels
 - An extra set of Sample bottles
 - An extra set of sample bottles for a duplicate sample
 - Shipping labels
 - Packing material
2. **Call the laboratory prior to sampling** to make sure there will be someone at the laboratory to receive and process the samples within 6 hours of sampling.
 3. **Consult flight schedules** to make sure there will be a flight that can get the samples to the laboratory within 6 hours of sampling.
 4. **Calibrate** equipment to be used for in situ measurements.
 5. Write the beach sampling location on the bottle label and Beach Sampling Data Sheet.
 6. Put on clean waders and gloves. Wade into the water to a depth of approximately 3 feet. Try to avoid kicking up sediment or wait until any sediment that has been kicked up settles. Stand downstream of the water current and wait for sediment to clear.
 7. Remove the bottle cap just before collecting the sample. Protect the cap from contamination. Do not to touch the inside of the bottle, or the inside of the cap.
 8. Open the sampling bottle and hold onto the base with one hand. Plunge the top of the bottle downward into the water. Avoid introducing surface scum. Point the mouth of the bottle into the current. Hold the bottle about 1 foot below the water surface and tip it slightly upward to allow air to exit and the bottle to fill.
 9. Remove the bottle from the water. Pour out a little water to leave airspace at the top of the jar. Fill two 250-mL bottles at each sampling location.
 10. Tightly close each bottle.
 11. Collect in situ field measurements using a handheld probe or similar. Collect in situ samples Immediately after collecting grab samples. Face upstream or into the current, allow any disturbed sediment to settle before submerging the probe to the manufacturer's suggested depth. Swirl the probe gently to allow good contact with the sensors. Wait for numbers to stabilize. Record results on field datasheets. Note that handheld probes must be calibrated prior to use in the field.

Collect replicates for half of the samples collected during the sampling season. To collect a replicate sample, you must first have requested extra jars from the laboratory. Repeat Steps 2 through 8 at the same location.

1. Complete bottle labels and attach them to each sample jar. Labels should be clean, waterproof, non-smearing, and large enough for all the information. Information on the label should include:

- Sample identifier (e.g., “city-date-sample” = “HOM-051507-01”)
 - Sample location (e.g., beach name)
 - Sampling date and time
 - Name of sampler
1. Wash your hands and arms with soap and water or waterless antimicrobial cleanser, or disinfectant lotion to reduce exposure to potentially harmful bacteria or microorganisms.

B.1.2 Sample Handling

Sample handling involves packing the samples in a cooler and shipping them to the laboratory. After sample collection is complete the samples must be handled with care so that they arrive to the laboratory in good condition. Step-by-step sample handling instructions are provided below:

2. Place the sample(s) in a pre-chilled cooler containing artificial ice to maintain a temperature from 1° to 10°C. Ask the laboratory ahead of time how much ice will be needed. Do not allow the samples to freeze.
3. Place enough packing material inside the cooler to protect the sample jars from breaking during transport to the laboratory.
4. Complete the chain-of-custody form. Put the form in a plastic bag and tape it to the inside of the cooler lid.
5. Write a note in the “Special Instructions” box requesting that the laboratory results be sent without delay (within 36 hours of sampling) to three people: the DEC BEACH Project Manager, the DEC BEACH Quality Assurance Officer, and you.
6. Fill out two custody seals and attach one to the front and one to the back of the cooler to span the lid seam. You want them to tear when the cooler is opened.
7. Securely tape the cooler shut prior to shipment. Attach shipping labels that identify the shipping destination and say: “keep cool,” “do not freeze,” and “fragile.”
8. Ship the samples to (Laboratory Name and Phone Number)

Remember that samples must be collected, shipped, and received by the laboratory in 6 hours.

Samples that exceed the 6-hour holding time may not be analyzed. Consult flight schedules and call the laboratory prior to sampling to make sure there will be a flight that can get the samples to the laboratory within 6 hours of sampling, and that there will be someone at the laboratory to receive the samples.

C. Appendix C: Example Chain of Custody Form



Admiralty Environmental
 641 W. Willoughby Ave., Suite 301
 Juneau, AK 99801
 (907) 483-4415

CHAIN OF CUSTODY/TRANSMITTAL RECORD
 PAGE 1 OF 1

CLIENT: Southeast Alaska Watershed Coalition				PROJECT: Hoonah Indian Association				AE 27176							
REPORT TO: Rebecca Bellmore				PHONE#: o: 907-205-4028 x3 c: 812-603-4910								# OF BOTTLES Fecal coliform, Enterococci			
ADDRESS: 1107 W. 8th St. Ste. #4 Juneau, AK 99801				SAMPLED BY: Jeremy Grant											
EMAIL: rebecca@sawcak.org				COMMENTS: Per client QAPP requirements, observe a 6 hour holding time. Rush data please.								FIELD RESULTS			
DATE	TIME	SITE DESCRIPTION / IDENTIFIER	MATRIX								pH	Temp °C			
8-3-2021	09:10	Curtina Creek #1	H ₂ O	1	1										
8-3-2021	09:11	Curtina Creek #1 dup	H ₂ O	1	1										
8-3-2021	09:20	Inner Point Sophia #2	H ₂ O	1	1										
RELINQUISHED BY:		RECEIVED BY:		RELINQUISHED BY:		RECEIVED BY:		Section to Be Completed by Receiving Laboratory							
Signature <i>[Signature]</i>		Signature <i>[Signature]</i>		Signature <i>[Signature]</i>		Signature <i>[Signature]</i>									
Printed Name Jeremy Grant		Printed Name Tim O'Neill		Printed Name		Printed Name									
DATE 8-3-2021		DATE 8/3/21		DATE		DATE									
TIME 09:18		TIME 1315		TIME		TIME									
								Temp °C: 7.97							
								Thermo ID#: Lab 7							
								Condition of Custody Seals: ✓							
								Initiated By: <i>[Signature]</i>							
								Shipped Via: _____							

D. Appendix E: Site-Specific Field Data Sheet

Date _____ Surveyor Name(s) _____

Site _____ Latitude _____ Longitude _____

Samples collected for (circle):

Fecal coliform Enterococci Fecal duplicate Enterococci duplicate MST

Field Measurements

Water Temperature _____ C°/F° Air Temperature _____ C°/F°

pH _____ Turbidity _____ (NTU)

Water clarity: Clear Murky Colored Other (describe)

Water smell: None Septic Algae Other (describe)

Wave height: Calm < 1 ft > 1 ft

Weather Conditions sunny partly sunny cloudy light rain rain heavy rain Other

Debris: in water? _____ (describe) _____
 on beach? _____ (describe) _____

# waterfowl	# dogs	# recreators		# boats
		on beach	in water	

Other Comments/observations:

Weather Station/App Records

Precipitation

Past 24 hr (in)	Past 48 hr (in)	Past 72 hr (in)	If past 74 hr = 0, most recent precip > 72 hr (date, amount)

Wind speed _____ (kts) Wind direction _____

Tide: High Low Ebb Flood