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2023 Cyanobacteria and Cyanotoxin Monitoring Work Plan in Selected Lakes in Indiana

PREPARED BY

Michelle Ruan Targeted Monitoring Section

Indiana Department of Environmental Management Watershed Assessment and Planning Branch Office of Water Quality 100 North Senate Avenue MC65-40-2 Shadeland Indianapolis, Indiana 46204-2251

April 28, 2023

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Approval Signatures

Ali Meils, Section Chief Targeted Monitoring Section

Vimolly Bownen

Timothy Bowren, Project Quality Assurance Officer Technical and Logistical Services Section

6/27/2023

6/28/2023

Date

Call Rel

Caleb Rennaker, Section Chief, Quality Assurance Manager Technical and Logistical Services Section

Kristen Arnold, Branch Chief, Quality Assurance Coordinator Watershed Assessment and Planning Branch

6/28/23

Date

6/27/23

Date

IDEM Quality Assurance staff reviewed and approves this work plan.

atrick Colos

Quality Assurance Staff IDEM Office of Program Support

6/29/23

Date

2023 Cyanobacteria and Cyanotoxin Monitoring Work Plan for Selected Lakes in Indiana B-057-OWQ-WAP-TGM-23-W-R0 April 28, 2023

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Work Plan Organization

This work plan is an extension of the existing Indiana Department of Environmental Management (IDEM) Office of Water Quality (OWQ) Watershed Assessment and Planning Branch (WAPB) March 2017 Quality Assurance Project Plan (QAPP) for Indiana Surface Water Programs (Surface Water QAPP) and Total Maximum Daily Limit (TMDL) program. The work plan serves as a link to the existing QAPP as well as an independent QAPP of the project. As per the U.S. EPA Guidance for Quality Assurance Project Plans, this work plan establishes criteria and specifications pertaining to a specific water quality monitoring project usually described in the following four groups and elements of a QAPP:

Group A. Project Management Planning

- A.1. Title and Approval Sheet
- A.2 Table of Contents
- A.3. Distribution List
- A.4. Project Organization
- A.5. Problem Definition and Background
- A.6. Project Description
- A.7. Quality Objectives and Criteria for Measurement Data
- A.8. Special Training and Certification
- A.9. Documents and Records

Group B. Data Generation and Acquisition

- B.1. Sampling Process Design (Experimental Design)
- B.2. Sampling Methods
- B.3. Sample Handling and Custody
- B.4. Analytical Methods
- B.5. Quality Control
- B.6. Instrument or Equipment Testing, Inspection, and Maintenance
- B.7. Instrument or Equipment Calibration and Frequency
- B.8. Inspection and Acceptance of Supplies and Consumables
- B.9. Nondirect Measurements
- B.10. Data Management

Group C. Assessment and Oversight

- C.1. Assessments and Response Actions
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Group D. Data Validation and Usability

- D.1. Data Review, Verification, and Validation
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- D.3. Reconciliation with User Requirements

2023 Cyanobacteria and Cyanotoxin Monitoring Work Plan for Selected Lakes in Indiana B-057-OWQ-WAP-TGM-23-W-R0 April 28, 2023

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List of Acronyms

ADDA	4E,6E-3-amino-9-methoxy-2,6,8-trimethyl-10-phenyldeca-4,6-dienoic acid
AIMS	Assessment Information Management System
CFR	Code of Federal Regulations
CLP	Contract Laboratory Program
CRQL	Contract Required Quantification Limit
CWA	Clean Water Act
CAAS	Cyanotoxin Automated Assay System
DO	Dissolved oxygen
DQA	Data quality assessment
ELISA	Enzyme-linked immunosorbant assay
GPS	Global Positioning System
HUC	Hydrologic Unit Code
IAC	Indiana Administrative Code
IC	Indiana Code
IDNR	Indiana Department of Natural Resources
MDL	Method detection limit
mg/L	Milligram per liter
µg/l	Micrograms per liter
µmho/cm	Micromho per centimeter
ml	Milliliter
NHD	National Hydrography Dataset
NTU	Nephelometric Turbidity Unit(s)
QA	Quality assurance
QAC	Quality assurance coordinator
QAM	Quality assurance manager
QAO	Quality assurance officer
QA/QC	Quality assurance and quality control
QC	Quality control
QAPP	Quality assurance project plan
RL	Reporting limit
RPD	Relative percent difference
SM	Standard methods
SS	Stainless steel
SU	Standard units
U.S.	United States
U.S. EPA	United States Environmental Protection Agency
WHO	World Health Organization

Program Objective

The IDEM OWQ is responsible for sampling and assessing Indiana's surface water pursuant to the Clean Water Act Section 305(b). IDEM's development of a monitoring program for lakes provides warnings to the public when cyanobacteria are present in large enough quantities and cyanotoxins are in high enough concentrations to render recreational water contact unsafe.

A. Project Management

A.1. Project Organization and Schedule

(QAPP Element A4)

The IDEM WAPB staff will sample designated Indiana Department of Natural Resources (IDNR) managed swimming beaches on lakes. Staff will evaluate samples for cyanobacteria cell counts and concentrations of the cyanotoxins microcystins, cylindrospermopsin (if producing species are present), anatoxin-a, and saxitoxin during the summer recreational season (May through August). Sample collection may occur outside of this timeframe at the request of IDNR. Collection of water chemistry samples will also take place for the following nutrient parameters: ammonia nitrogen, nitrate plus nitrite, total Kjeldahl nitrogen, and total phosphorous.

Sampling will take place at the following reservoirs or lakes: Worster Lake at Potato Creek State Park, Sand Lake at Chain O'Lakes State Park, Lake James at Pokagon State Park, Lake James at Pokagon State Park Inn, Salamonie Reservoir at Lost Bridge West State Recreation Area, Mississinewa Reservoir at Miami State Recreation Area, Cecil M. Harden Lake at Raccoon State Recreation Area, Monroe Lake at Paynetown and Fairfax State Recreation Areas, Hardy Lake at Hardy Lake State Recreation Area, Whitewater Lake at Whitewater Memorial State Park, Deam Lake State Recreation Area, Whitewater Lake at Whitewater Starve-Hollow State Recreation Area, Brookville Reservoir at Mounds and Quakertown State Recreation Areas, Summit Lake at Summit Lake State Park, Cagles Mill Lake at Lieber State Recreation Area, Kunkel Lake at Ouabache State Park, Ferdinand State Forest Lake at Ferdinand State Forest, Lake Lincoln at Lincoln State Park, and Patoka Lake at Newton-Stewart State Recreation Area.

Sampling will occur at selected sites on these lakes monthly unless cyanobacteria cell counts exceed 100,000 cells/ml. Excessive cell counts will initiate biweekly sampling. If a high cell count is due to picocyanobacteria and picocyanobacterial cells comprise 80% of the sample, a follow-up visit is not necessary. Toxin concentrations that exceed the Human Recreation Advisory thresholds in Table 1 will initiate weekly sampling. Additionally, sampling for field parameters and cyanotoxin analysis will happen at the Dog Park Lake at Fort Harrison State Park before it opens on March 1st, monthly from May through June, and then biweekly from July 1st through October 31st, or until the lake closes. The addition of sites and analyses may occur depending on time and staffing constraints.

A.2. Background and Project Description

(QAPP Elements A5, A6)

The presence of sufficient numbers of cyanobacteria in surface waters serves as an indicator of possible cyanotoxin production. Exposure to high counts of cyanobacterial cells and high cyanotoxin concentrations may lead to various illnesses in humans and animals. Exposure is through contact with or ingestion of surface water. Indiana has yet to establish cyanobacteria or cyanotoxin water quality criteria or standards for full body contact recreational uses to protect human health. However, the World Health Organization (WHO) guidelines are in common use for this purpose. The United States Environmental Protection Agency (U.S. EPA) recently published recreation guidelines. Indiana will use a combination of WHO and U.S. EPA recommended values described in Table 1. Dog Park Lake values are based on suggestions from the California Environmental Protection Agency's Office of Environmental Health Hazard Assessment and the Oregon Health Authority.

Cyanotoxins will be analyzed using Abraxis[™] enzyme linked immunosorbant assay (ELISA) test kits (microcystins, cylindrospermopsin, anatoxin-a, and saxitoxin) in the IDEM WAPB Algae Laboratory. The Cyanotoxin Automated Assay System (CAAS) performs automated sample processing and reading. If samples need manual processing, samples will be read with the Chromate Reader. In addition to cyanobacteria sampling, water chemistry samples will be collected for ammonia nitrogen, nitrate plus nitrite as nitrogen, total Kjeldahl nitrogen, and total phosphorous. Staff will record the following field parameters: pH, DO, DO percent saturation, temperature, turbidity, and specific conductivity. Observations of atmospheric and lake or beach conditions will also be recorded for each sampling event. At the Dog Park Lake, staff will record field measurements and collect grab samples for cyanotoxin analysis. For Ferdinand State Forest Lake, Lake Lincoln, and Patoka Lake, IDNR staff will collect cyanobacteria and cyanotoxin samples and deliver samples to the IDEM WAPB Algae Laboratory. Staff will enter all results into the WAPB Assessment Information Management System (AIMS) database and post results on <u>www.algae.IN.gov</u>.

A.3. Quality Objectives and Criteria

(QAPP Element A7)

Guidance on Systematic Planning Using the Data Quality Objective Process (U.S. EPA 2006) describes a planning tool for environmental data collection activities. The process provides a basis for balancing decision uncertainty with available resources. This work plan contains the products of the process which include the data quality objectives and measurement quality objectives. U.S. EPA recommends the Data Quality Objective Process for all significant environmental data collection projects. The process is a seven-step systematic planning process used to clarify study objectives, define the appropriate types of data, and establish decision criteria on which to base the final use of the data. Numbers 1. through 7. outline the data quality objectives process outputs.

1. State the Problem

The waters in the previously identified recreational lakes either have a documented history of or the potential of supporting cyanobacteria blooms which may present a human and animal health hazard.

2. Identify the Goals of the Study

This project will provide information regarding the nature and extent of cyanobacteria populations in the selected swimming areas, and will measure microcystins, cylindrospermopsin, anatoxin-a, and saxitoxin concentrations. By routine monitoring of cyanobacteria and cyanotoxins in lakes, IDEM determines whether conditions are present to pose an unreasonable health risk to swimmers or dogs. Steps are taken to adequately warn the public of potential risks.

3. Identify Information Inputs

Collect composite water samples at each swimming beach's sampling locations for cyanobacteria identification and enumeration, and cyanotoxin analyses. Identify and enumerate cyanobacterial cells in each sample using microscopy. Analysis for each cyanotoxin requires a specific immunoassay test kit: Abraxis[™] ADDA ELISA for microcystins; and Abraxis[™] ELISA for cylindrospermopsin, anatoxin-a, and saxitoxin.

Measure field parameters (pH, DO, DO percent saturation, temperature, turbidity, and specific conductance) at each site during each sampling event.

Collect water samples for ammonia nitrogen, nitrate and nitrite, total Kjeldahl nitrogen, and total phosphorous.

Include visual field observations, weather conditions, and lake or beach conditions.

For the Dog Park Lake, only take field measurements and collect cyanotoxin grab samples.

4. Define the Boundaries of the Study Area

The study areas consist of the designated swimming beaches at Worster Lake at Potato Creek State Park, Sand Lake at Chain O'Lakes State Park, Lake James at Pokagon State Park, Lake James at Pokagon State Park Inn, Salamonie Reservoir at Lost Bridge West State Recreation Area, Mississinewa Reservoir at Miami State Recreation Area, Cecil M. Harden Lake at Raccoon State Recreation Area, Monroe Lake at Paynetown and Fairfax State Recreation Areas, Hardy Lake at Hardy Lake State Recreation Area, Whitewater Lake at Whitewater Memorial State Park, Deam Lake at Deam Lake State Recreation Area, Starve-Hollow Lake at Starve-Hollow State Recreation Area, Brookville Reservoir at Mounds and Quakertown State Recreation Areas, Summit Lake at Summit Lake State Park, Cagles Mill Lake at Lieber State Recreation Area, Kunkel Lake at Ouabache State Park, the Dog Park Lake at Fort Harrison State Park, Ferdinand State Forest Lake at Ferdinand State Forest, Lake Lincoln at Lincoln State Park, and Patoka Lake at Newton-Stewart State Recreation Area. Sampling sites are listed in Table 2 and can be viewed in Figures 1 through 20.

Sites will be sampled monthly from May through August. Excessive cell counts (≥100,000 cells/ml) will initiate biweekly sampling. Toxin concentrations that exceed the Human Recreation Advisory thresholds in Table 1 will initiate weekly sampling. The Dog Park Lake at Fort Harrison State Park will be sampled before it opens on March 1st and monthly from May through June. Biweekly sampling will start from July 1st through October 31st, or until the lake closes. Detection of any toxin will initiate weekly sampling.

Sampling activities will not be conducted during hazardous weather conditions (e.g., thunderstorms or heavy rain in the vicinity) or when unexpected physical barriers to accessing the site exist. In the case of hazardous weather conditions, staff may wait until it is safe to enter the water to collect samples. If the beach at the sampling location is flooded, staff will collect samples from where they are able to and make notations on the field sampling sheet. The field crew chief will make the final determination as to whether or not a site is safe to enter.

5. Develop the Analytic Approach

Compare cyanobacteria cell counts to the WHO and U.S. EPA guidelines. IDEM will post recreational advisory signs when the cyanobacteria cell counts are at or above 100,000 cells/ml.

Table 1 summarizes the cyanotoxin exposure thresholds. U.S. EPA guidelines recommend posting recreational advisories when microcystin concentrations are at or above 8 µg/l and IDNR issues an additional caution advisory indicating children and the immunocompromised should not swim (U.S. EPA 2019). WHO guidelines recommend prohibiting swimming when microcystin concentrations are at or above 20 µg/l. IDNR will use the WHO recreational guideline value of 6 µg/l for a caution advisory for cylindrospermopsin concentrations and U.S. EPA recommended cylindrospermopsin limit of 15 µg/l for beach closures. Using the State of Ohio recommended thresholds for anatoxin-a and saxitoxin, IDNR will post a caution advisory when concentrations are at or above 8 µg/l and 0.8 µg/l respectively (State of Ohio 2020). The California Environmental Protection Agency's Office of Environmental Health Hazard Assessment and the Oregon Health Authority action levels are used for microcystin, anatoxin-a, cylindrospermopsin, and saxitoxin exposure for dogs. Swimming is prohibited at the Dog Park Lake when concentrations are at or above 0.8 µg/l microcystin, 1.0 µg/l of cylindrospermopsin, 0.4 µg/l anatoxin-a, and 0.05 µg/l saxitoxin (Linville 2017). An advisory will be issued if microcystin is detected at or above 0.4 µg/L or cylindrospermopsin is detected at or above 0.5 µg/l, which is half of the respective closing limit.

Exposure Reference Values (µg/l)	Microcystin	Cylindrospermopsin	Anatoxin-a	Saxitoxin	
Human Recreation Advisory	8	6	8	0.8	
Human Recreation Prohibited	20	15	30	3	
Dog Recreation Advisory	0.4	0.5	_	-	
Dog Recreation Prohibited	0.8	1.0	0.4	0.05	
The reporting limits for anatoxin-a and saxitoxin are the same as the respective closure thresholds.					

Table 1. Exposure Thresholds

6. Specify Performance or Acceptance Criteria

Good quality data are essential for minimizing decision error. By minimizing both sampling design error and laboratory error, the public can place more confidence in the cell counts and cyanotoxin concentrations reported through this program. Use program-specific controls to minimize the introduction of errors. Controls for field sampling include the collection of field duplicates and blanks, and field audits. Laboratory controls include thorough verification of species identifications and cyanotoxin analyses. Take and store photographs of cyanobacterial identifications. Staff will be trained on how to collect samples, operate field and lab equipment, and perform required quality assurance and quality control (QA/QC) procedures. QA/QC review reports are developed as a final step of QA process and data assessment. The data and their usability identified in the QA/QC review reports are used in the decision making for water quality assessments.

7. Develop the Plan for Obtaining Data

Composite sampling at each beach swimming area ensures a better representation of the cyanobacteria populations in the swimming area. Staff will wade in to about waist high and stand for a minute to let the water and sediment settle before taking the sample using the integrated water column sampler (IDEM 2023a). Collect samples in the morning and afternoon when cyanobacteria are believed to be the most prevalent. At the Fort Harrison Dog Park Lake, collecting samples near the shore reflects dogs' drinking habits.

A.4. Specialized Training or Certification and Staffing Requirements

(QAPP Element A8)

Written standard operating procedures (SOPs) are available to WAPB staff. Experienced peers train staff on how to collect surface water samples, operate field and lab equipment, and perform required quality assurance and quality control (QA/QC) procedures. Staff training and SOP techniques are based on proven U.S. EPA sampling methods and recommendations for containers, preservation techniques, and holding times as cited in the Code of Federal Regulations (CFR) (CFR 2023).

Assign two staff to each field sampling run. If necessary, one staff can sample. Senior staff will provide training and review for IDEM staff and summer interns on how to collect surface water samples, operate field and lab equipment, and perform QA/QC procedures.

Select WAPB staff received training in algal identification and enumeration from staff at PhycoTech in St. Joseph, Michigan. A representative from Eurofins Abraxis[™] trained select WAPB staff on the use of the Eurofins Abraxis[™] CAAS instrument. Staff trained in cyanobacteria identification and enumeration, and use of the Eurofins Abraxis[™] CAAS will perform these tasks in the IDEM WAPB Algae Laboratory.

Targeted Monitoring, Technical and Logistical Services, and Probabilistic Monitoring Sections staff share data handling. Sampling staff will input field observations and field data into the AIMS database. Sampling staff will perform QA/QC of all field data entry. Chemists from the Technical and Logistical Services Section will validate the data through QA/QC procedures cited in Section D of this work plan.

B. Data Generation and Acquisition

B.1. Sampling Process Design (Experimental Design)

(QAPP Element B1)

The Targeted Monitoring Section chief will select the initial sample site locations based on the availability of a public swimming beach at a state or federally managed recreation area. Table 2 lists the selected sampling sites with associated lake name, AIMS L-site, county name, HUC12, and latitude and longitude of each sampling site. Figures 1 through 20 depict all sampling sites for each lake. The Trimble Juno[™] Series Global Positioning System, with an accuracy of 1-3 meters, determined the final coordinates for each site during the project's presurvey phase.

Site #	Lake Name	AIMS L-site	County	HUC12	Latitude	Longitude
1	Whitewater	GMW-07-0001	Union	050800030712	39° 36' 54.9"	-84° 57' 58.1"
2	Hardy	WEM-06-0001	Scott	051202070604	38° 46' 55.2"	-85° 41' 53.8"
3	Cecil M Harden	WLV-12-0002	Parke	051201081208	39° 44' 48.9"	-87° 04' 33.5"
4	Monroe - Paynetown	WEL-07-0001	Monroe	051202080702	39° 04' 49.1"	-86° 25' 51.1"
5	Monroe - Fairfax	WEL-07-0003	Monroe	051202080703	39° 01' 20.5"	-86° 28' 53.9"
6	Mississinewa	WMI-06-0001	Wabash	051201030605	40° 42' 08.9"	-85° 56' 08.0"
7	Worster	UMK-01-0005	St Joseph	071200010104	41° 33' 03.2"	-86° 21' 41.8"
8	Salamonie	WSA-04-0001	Huntington	051201020405	40° 46' 46.1"	-85° 38' 14.9"
9	Sand	LMJ-16-0024	Noble	040500011601	41° 19' 50.0"	-85° 23' 00.9"
10	Lake James - Pokagon SP	LMJ-08-0033	Steuben	040500010803	41° 42' 35.5"	-85° 02' 10.7"

Table 2. Sampling Sites

2023 Cyanobacteria and Cyanotoxin Monitoring Work Plan for Selected Lakes in Indiana B-057-OWQ-WAP-TGM-23-W-R0 April 28, 2023

Site #	Lake Name	AIMS L-site	County	HUC12	Latitude	Longitude
11	Lake James - Potawatomi Inn	LMJ-08-0078	Steuben	040500010803	41° 42' 09.3"	-85° 01' 21.7"
12	Brookville - Quakertown	GMW-07-0005	Union	050800030715	39° 35' 10.9"	-84° 59' 41.5"
13	Brookville - Mounds	GMW-07-0004	Franklin	050800030717	39° 29' 53.2"	-84° 59' 12.6"
14	Deam	OSK-07-0001	Clark	051401010702	38° 27' 58.5''	-85° 51' 40.8''
15	Starve-Hollow	WEM-09-0007	Jackson	051202070905	38° 48' 50.4''	-86° 04' 50.3''
16	Kunkel	WUW-06-0005	Wells	051201010604	40° 43' 12.4"	-85° 06' 21.0"
17	Fort Harrison Dog Park	WWU-09-0027	Marion	051202010904	39° 52' 39.4''	-86° 01' 02.5"
18	Summit	WED-01-0001	Henry	051202040102	40° 01' 33.0"	-85° 18' 28.6"
19	Cagles Mill	WWE-05-0015	Putnam	051202030512	39° 28' 29.1"	-86° 53' 54.5"
20	Ferdinand State Forest	OLP-04-0010	Dubois	051402010402	38° 15' 21.8"	-86° 46' 35.1"
21	Lake Lincoln	OLP-09-0012	Spencer	051402010905	38° 06' 13.2"	-86° 59' 41.4"
22	Patoka	WPA-01-0016	Orange	051202090106	38° 24' 50.5"	-86° 40' 4.4"









Figure 3. Hardy Lake Site Map





Figure 4. Cecil M. Harden Lake Site Map



Figure 5. Monroe Lake Site Map



Figure 6. Mississinewa Lake Site Map



Figure 7. Worster Lake Site Map



Figure 8. Salamonie Lake Site Map



Figure 9. Sand Lake Site Map

Figure 10. Lake James Site Map





Figure 11. Brookville Lake Site Map

Figure 12. Deam Lake Site Map





Figure 13. Starve Hollow Lake Site Map







Figure 15. Fort Harrison Dog Park Lake Site Map



Figure 16. Summit Lake Site Map



Figure 17. Cagles Mill Lake Site Map







Figure 19. Lake Lincoln Site Map

Figure 20. Patoka Lake Site Map



B.2. Sampling Methods and Sample Handling

(QAPP Elements B2, B3)

1. Collecting Surface Water Samples for Cyanobacteria and Cyanotoxin Analysis

Staff will wade into the water body to a level about waist high and stand still for a minute to let the water and sediment settle before taking a sample using the integrated water column sampler. Collect samples from three locations (each side and the middle portion) at each beach swimming area. Combine the samples in the churn to form a composite sample. Take samples for cyanobacteria identification and enumeration, and cyanotoxin analysis from the composite sample. Details of sampling equipment and methods are in the Collecting Surface Water Samples for Cyanobacteria and Cyanotoxin Analysis TSOP (IDEM 2023a).

At the Fort Harrison Dog Park Lake, collect grab samples near the shore to reflect dogs' drinking habits. Staff will check the pH of the buffered cyanotoxin vial to make sure it is between 6 to 8 due to sample storage requirements for anatoxin-a.

2. Water Chemistry Sampling

Staff will collect nutrient samples, record water chemistry field measurements, and note lake or beach conditions on the IDEM Stream Sampling Field Data Sheet. Water chemistry sampling will adhere to the Water Chemistry Field Sampling Procedures TSOP (IDEM 2020a). Collection of nutrient samples will diverge from the TSOP in that the nutrient sample bottles will be filled from the composite sample in the churn.

3. Field Parameter Measurements

Measure DO, pH, water temperature, specific conductance, and DO percent saturation with a data sonde each time water samples are collected. Measure field parameters directly from the lake water.

B.3. Analytical Methods

(QAPP Element B4)

- 1. Laboratory Procedure for Cyanobacteria Identification and Enumeration Cyanobacteria cells are identified and quantified for each sample using a microscope with brightfield Nomarski differential interference contrast and epifluorescence capabilities. These processes are detailed in the Cyanobacteria Identification and Enumeration TSOP (IDEM 2023b).
- 2. Laboratory Procedure for Microcystins, Cylindrospermopsin, Anatoxin-a, and Saxitoxin Analysis

Samples are analyzed for microcystins, cylindrospermopsin, anatoxin-a, and saxitoxin concentrations using Abraxis[™] ELISA test kits. The Determination of Cyanobacteria Toxins in Ambient and Drinking Water by ELISA TSOP contains procedures for sample preparation and analysis for each cyanotoxin (IDEM 2021a).

3. Nutrient Parameters Measurements

Pace Analytical Services, Inc. (Indianapolis, Indiana) will process nutrient samples in accordance with preapproved test methods and within allotted time frames. A chain-of-custody form accompanies each sample set through the analytical process. Table 3 identifies the nutrient parameters, respective test methods, and quantification limits.

Parameter	Method	Limits of Quantification	Units	Preservative	Holding Times
Ammonia Nitrogen	U.S. EPA 350.1	0.1	mg/L	H ₂ SO ₄ < pH 2	28 days
Nitrogen, Nitrate and Nitrite	U.S. EPA 353.2	0.1	mg/L	H ₂ SO ₄ < pH 2	28 days
Total Kjeldahl Nitrogen (TKN)	U.S. EPA 351.2	0.5	mg/L	H ₂ SO ₄ < pH 2	28 days
Total Phosphorus	U.S. EPA 365.1	0.05	mg/L	H ₂ SO ₄ < pH 2	28 days

Table 3. Nutrient Parameters Test Methods

4. Field Parameter Measurements

Field measurements of DO, temperature, pH, conductivity, and turbidity will be taken each time a water sample is collected. Table 4 identifies the field parameters to measure at each sampling site, respective test methods, and quantification limits. Document field observations from each site on field sheets during each sampling run.

Parameter	Method	Limits of Quantitation	Units
DO (data sonde optical)	ASTM D888-09 ¹	0.05	mg/L
DO (membrane probe)	SM 4500-OG ¹	0.05	mg/L
DO % Saturation (data sonde optical)	ASTM D888-09	0.05	%
DO % Saturation (membrane probe)	SM 4500-OG	0.01	%
Turbidity (Hach™ turbidity kit)	U.S. EPA 180.1 ¹	0.05	NTU
Specific Conductance (data sonde)	SM 2510B	1.00	µmho/cm
Temperature (data sonde)	SM 2550B(2)	0.1	°Celsius
Temperature (field meter)	SM 2550B(2) ¹	0.1	°Celsius
pH (data sonde)	U.S. EPA 150.2	0.10	SU
pH (field meter)	SM 4500H-B ¹	0.10	SU
¹ Method used for field calibration ch	eck		

 Table 4. Field Parameters Test Methods

B.4. Quality Control and Custody Requirements

(QAPP Elements B5, B3)

1. Cyanobacteria Cell Count Data

AIMS will assign quality control (QC) samples for each sampling trip prior to sample collection. This includes the field blank, field duplicate, and matrix spike/matrix spike

duplicate samples. IDEM taxonomists will follow procedures outlined in the Cyanobacteria Identification and Enumeration TSOP (IDEM 2023b). As with normal samples, identify and quantify cyanobacteria cells for each QC sample using a microscope with brightfield Nomarski differential interference contrast and epifluorescence capabilities.

Rinse the nannoplankton counting chamber and coverslip thoroughly with deionized water and dry to avoid cross-contamination between samples. Analyze one field blank sample each week to ensure samples are not contaminated in the field or laboratory. This also serves as a check to ensure the coverslip and counting chamber are adequately rinsed between each sample. Analyze one field duplicate sample each week. The relative percent difference (RPD) between normal and field duplicate cell counts should be at or below 20%. The RPD between IDEM taxonomists should also be at or below 20%.

2. Eurofins Abraxis[™] ELISA

Analytical results from the Eurofins Abraxis[™] ELISA include QC checks, which can determine precision, accuracy, and completeness for each batch of samples. Archive raw data by analytical batch for easy retrieval and review. Follow chain-of-custody procedures, including date and time of sample collection; time of sample set-up for cyanotoxin analysis and the identification process; time of reading the results; and time and method of disposal. Thoroughly document any method deviations in the raw data bench sheets and records. Refer to the Determination of Cyanobacteria Toxins in Ambient and Drinking Water by ELISA TSOP for QA/QC protocols (IDEM 2021a). Collect all QA/QC samples according to the following guidelines:

Field Duplicate	1 per batch or at least 1 for every 20 samples collected (≥ 5%)
Field Blank	1 per batch or at least 1 for every 20 samples collected ($\geq 5\%$)
Laboratory Blank	At least 1 laboratory blank (sterile laboratory water blanks) per analysis batch, per day, or every time the laboratory is in use
Laboratory Fortified Blank	At least 1 per analysis batch (spike solution added to sterile laboratory water blanks)
Matrix Spike/Matrix Spike Duplicate	At least 1 per analysis batch (spike solution added to field samples) Should a matrix spike value exceed the calibration range, analyze a laboratory duplicate
	instead.

3. Nutrient Data

AIMS generates a quality control (QC) sample collection schedule containing field blanks, equipment blanks, and duplicate samples for each sampling project. Use sample bottles and preservatives certified for purity. Sample collection containers for each parameter, preservative, and holding time will adhere to U.S. EPA requirements. Collect field duplicates, and matrix spike and matrix spike duplicates (MS/MSD) at the rate of one per sample analysis set or one per every 20 samples, whichever is greater. Take field blank samples, using ASTM D1193-91 Type I water, at a rate of one per sample analysis set or one per every 20 samples, whichever is greater.

For each sampling event, fill out a Chain-of-Custody Form which will accompany the sample bottles from sample collection until delivery to the contract lab. Water Chemistry Field Sampling Procedures TSOP (IDEM 2020a) describes sampling procedures in more detail. QA documentation for each batch of samples consists of a Chain-of-Custody Form, bench sheets, spreadsheets of results, and the QC report generated by staff from the Technical and Logistical Services Section.

B.5. Field Instrument Testing and Calibration

(QAPP Elements B6, B7)

This project will benefit from the planned upkeep of the measurement instruments, which enhances the instrument performance, and ensures accurate and precise readings. Maintain equipment logs which record equipment calibration and status.

Measurement equipment requires periodic calibration or standardization to reliably produce accurate results. Calibrate the multiparameter data sonde immediately prior to each week's sampling (IDEM 2020a; IDEM 2020b). Record, maintain, store, and archive calibration results and drift values in logbooks located in the WAPB calibration laboratories. The drift value is the difference between two successive calibrations. Field parameter calibrations will conform to the procedures as described in the instrument user manuals (IDEM 2020b). Conduct the DO component of the calibration procedure using the air calibration method. When used in conjunction with bacteriological and water chemistry sampling, field check the unit for accuracy once during the week by comparison with a YSI EcoSense DO200A DO meter or YSI ProSolo ODO meter, as well as with a Hach™ turbidimeter and an Oakton pH meter (IDEM 2020a, pages 24-26). Record field calibration values on field sheets and enter into the AIMS database. Also conduct a DO meter reading at sites where the DO concentration is 4.0 mg/L or less. Calibration of YSI Multiparameter Data Sondes TSOP documents procedures for field instrument calibration and is followed by all staff participating in data collection for this study (IDEM 2020b).

C. Assessment and Oversight

(QAPP Elements C1, C2)

Conduct field and laboratory performance and system audits to ensure good quality data. The

field and laboratory performance includes precision measurements by relative percent difference of field and laboratory duplicates; accuracy measurements by percent of recovery of MS/MSD samples analyzed in the laboratory; and completeness measurements by the percent of planned samples actually collected, analyzed, reported, and usable for the project.

Field audits ensure sampling activities adhere to approved SOPs. WAPB management staff systematically conduct audits of all WAPB staff engaging in field sampling activities.

C.1. Data Quality Assessment Levels

The samples and various types of data collected by this program are intended to meet the QA criteria and data quality assessment (DQA) levels as described in the Surface Water QAPP (IDEM 2017, pages 181-183). Field data collected as specified in this work plan will comply with DQA Level 2 requirements. All field data are QA/QC'd for completeness, precision, and accuracy. Nutrient data submitted from Pace Analytical Services, Inc. (Indianapolis, Indiana) will adhere to DQA Level 3. Cyanotoxin data analyzed in IDEM WAPB Algae Laboratory are assigned DQA Level 4. QA/QC review reports are developed as a final step of QA process and data assessment. The data and their usability identified in the QA/QC review reports are used in the decision making for water quality assessments.

D. Data Validation and Usability

(QAPP Element D1, D2)

D.1. Quality Assurance, Data Qualifiers and Flags

The various data qualifiers and flags used for QA and validation of the data are found on pages 184-185 of the Surface Water QAPP (IDEM 2017).

D.2. Data Usability

The environmental data collected, and its usability are qualified and classified into one or more of the four Categories: Enforcement Capable, Acceptable Data, Estimated Data, or Rejected Data.

- 1. **Enforcement capable** results are DQA Level 3 or 4 data which meet all QC checks.
- 2. Acceptable data are DQA Level 2, 3, or 4 data suitable for decision making. Although a few data may be estimated or even unusable, the sample set, as a whole, has scientific and statistical integrity. Scientific and statistical decisions may be made with respect to the data quality objectives.
- 3. **Estimated data** may be suitable for enforcement or decision making on a caseby-case basis. Estimated data are suitable for determining future sampling needs, locating target parameters, and identifying possible contaminant levels.
- 4. Rejected data are not suitable for enforcement or for decision making.

D.3. Laboratory and Estimated Cost

This project will take a minimum of four sampling days (day trips) or a maximum of eight sampling days per month. If the cell count for a site exceeds 100,000 cells/ml, sampling frequency will increase to two weeks and if toxin concentrations exceed the Human Recreation Advisory thresholds, the site will be revisited weekly until concentrations fall below threshold levels. Sampling trips require one vehicle and two staff.

Laboratory analysis and data reporting for this project will comply with the Surface Water QAPP (IDEM 2017), Request for Proposals 21-68153 (IDEM 2021b), and the IDEM Quality Management Plan (IDEM 2018). IDEM WAPB Algae Laboratory staff will analyze cyanobacteria and cyanotoxin samples. Identification and enumeration of cyanobacteria samples is performed in-house and does not incur a cost per sample. Laboratory supplies to run cyanotoxin analyses using Abraxis[™] ELISA test kits will cost approximately \$33,101. Pace Analytical Services, Inc. (Indianapolis, Indiana) will perform tests on the nutrient parameters outlined in Table 3 for a total cost of \$19,156. This estimated lab cost is based on the number of samples analyzed in 2022, with a 10% contingency added for margin of error. The anticipated total budget for laboratory costs for this project is \$52,257.

D.4. Reference Manuals and Personnel Safety

Role	Required Training/Experience	Training References	Training Notes
All staff	-Basic first aid and	-A minimum of 4 hours	- WAPB staff meeting Health
participating in	cardiopulmonary	of in-service training	and Safety Training
field activities	resuscitation (CPR)	provided by WAPB (IDEM 2010)	requirements will accompany staff lacking 4 hours of in- service training or appropriate certification in the field at all times.
	-Personal Protective Equipment (PPE) Policy	-IDEM 2008	-When working on boundary waters as defined by Indiana Code (IC) <u>14-8-2-27</u> or between sunset and sunrise
	-Personal flotation devices (PFD)	-February 29, 2000, WAPB internal memorandum regarding use of approved PFDs	on any waters of the state, all personnel in the watercraft must wear a high intensity whistle and Safety of Life at Sea (SOLAS) certified strobe light.
All staff performing laboratory duties	-Review relevant safety procedures	-OWQ Laboratory Safety Plan (IDEM 2021c)	
	-Review safety data sheets annually	-IDEM Hazard Communication (HazCom) Plan (IDEM 2019)	

Table 5. Personnel Safety and Reference Manuals

References

- Code of Federal Regulations 2023. <u>40 CFR Part 136 Guidelines Establishing Test</u> <u>Procedures for the Analysis of Pollutants</u>. Title 40 was last amended March 29, 2023.
- U.S. EPA 2002. <u>Guidance for Quality Assurance Project Plans. EPA QA/G-5</u>. EPA/240/R-02/009. U.S. EPA, Office of Environmental Information, Washington, D.C.
- U.S. EPA 2006. <u>Guidance on Systematic Planning Using the Data Quality Objectives Process.</u> <u>EPA QA/G-4</u>. EPA/240/B-06/001. U.S. EPA, Office of Environmental Information, Washington, D.C.
- U.S. EPA 2019. <u>Recommended Human Health Recreational Ambient Water Quality Criteria or</u> <u>Swimming Advisories for Microcystins and Cylindrospermopsin</u>. EPA 822-R-19-001. U.S. EPA, Office of Water, Health and Ecological Criteria Division, Washington, D.C.
- Indiana Administrative Code, <u>Title 327 Water Pollution Control Division, Article 2. Water</u> <u>Quality Standards</u>. Last updated March 29, 2023. Available at <u>http://www.in.gov/legislative/iac/iac_title?iact=327</u>.
- IDEM 2008. <u>Personal Protective Equipment Policy</u>. A-059-OEA-08-P-R0. IDEM, Indianapolis, Indiana.
- IDEM 2010. <u>Health and Safety Training Policy</u>. A-030-OEA-10-P-R2. IDEM, Indianapolis, Indiana.
- IDEM 2017. <u>Quality Assurance Project Plan (QAPP) for Indiana Surface Water Programs</u>. B-001-OWQ-WAP-XX-17-Q-R4. OWQ, WAPB, Indianapolis, Indiana.
- IDEM 2018. IDEM Quality Management Plan 2018. IDEM, Indianapolis, Indiana.
- IDEM 2019. IDEM Hazard Communication (HazCom) Plan. IDEM, Indianapolis, Indiana.
- IDEM 2020a. <u>Water Chemistry Field Sampling Procedures</u>. B-015-OWQ-WAP-XXX-20-T-R0. OWQ, WAPB, Indianapolis, Indiana.
- IDEM 2020b. <u>Calibration of YSI Multiparameter Data Sondes</u>. B-014-OWQ-WAP-XXX-20-T-R0. OWQ, WAPB, Indianapolis, Indiana.
- IDEM 2021a. <u>Determination of Cyanobacteria Toxins in Ambient and Drinking Water by ELISA</u>. S-001-OWQ-WAP-TGM-21-T-R4. OWQ, WAPB, Indianapolis, Indiana.
- IDEM 2021b. State of Indiana Request for Proposals 21-68153, Solicitation for: Laboratory Analytical Services, Indiana Department of Administration, Indianapolis, Indiana.
- IDEM 2021c. <u>OWQ Watershed Assessment and Planning Branch Laboratory Safety Plan</u>. IDEM, Indianapolis, Indiana.
- IDEM 2023a. <u>Collecting Surface Water Samples for Cyanobacteria and Cyanotoxin Analysis</u>. S-003-OWQ-WAP-TGM-23-T-R1. OWQ, WAPB, Indianapolis, Indiana.

- IDEM 2023b. <u>Cyanobacteria Identification and Enumeration</u>. B-002-OWQ-WAP-XX-23-T-R1. OWQ, WAPB, Indianapolis, Indiana.
- State of Ohio 2020. <u>Harmful Algal Bloom Response Strategy for Recreational Waters</u>. Rev.8.4.2020.
- Clesceri, L.S., Greenburg, A.E., Eaton, A.D., 2017. SM-Standards Methods for the Examination of Water and Wastewater 23rd Edition. American Public Health Association.
- John, D.M., B.A. Whitton and A.J. Brook. 2002. The Freshwater Algal Flora of the British Isles: An Identification Guide to Freshwater and Terrestrial Algae. Cambridge University Press. New York. 702 pages.
- Linville, Regina, PhD, Toxicologist, Office of Environmental Health Hazard Assessment, California Environmental Protection Agency. Email correspondence, February 8, 2017.
- Prescott, G.W. 1978. How to Know the Freshwater Algae. Third Edition. Wm. C. Brown, Dubuque, IA.
- Prescott, G.W. 1982. Algae of the Western Great Lakes Area. Second Edition. Otto Koeltz Science Publishers, Koenigstein.
- Wehr, J.D. and R.G. Sheath. 2003. Freshwater Algae of North America. Academic Press, Boston. 918 pages.
- World Health Organization 2003. Guidelines for safe recreational waters Volume 1 Coastal and fresh waters.
- World Health Organization 2021. <u>Toxic Cyanobacteria in Water</u>. Second Edition. CRC Press, London. 858 pages.

Distribution List

Electronic Distribution Only (QAPP Element A3)

Name	Organization
Kristen Arnold	IDEM, OWQ, WAPB
Tim Bowren	IDEM, OWQ, WAPB, Technical and Logistical Services Section
Todd Davis	IDEM, OWQ, WAPB, Targeted Monitoring Section
Charlie Hostetter	IDEM, OWQ, WAPB, Technical and Logistical Services Section
David Jordan	IDEM, OWQ, WAPB, Technical and Logistical Services Section
Ali Meils	IDEM, OWQ, WAPB, Targeted Monitoring Section
Caleb Rennaker	IDEM, OWQ. WAPB, Technical and Logistical Services Section
Michelle Ruan	IDEM, OWQ, WAPB, Targeted Monitoring Section
Addison Seidler	IDEM, OWQ, WAPB, Targeted Monitoring Section
QA Staff	IDEM, OPS, Recycling Education and QA, QA Manager