HEE NERR Nutrient Metadata January-December 2022

Latest Update: June 14, 2023

Note: This is a provisional metadata document; it has not been authenticated as of its download date. Contents of this document are subject to change throughout the QAQC process and it should not be considered a final record of data documentation until that process is complete. Contact the CDMO (cdmosupport@baruch.sc.edu) or reserve with any additional questions.

I. Data Set and Research Descriptors

1) Principal investigator(s) and contact persons -

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2) Research objectives -

a) Monthly grab sampling program

Our long term monitoring program consists of 4 official SWMP sites in our watershed. The overall goal is to track the physical and biogeochemical parameters of the He'eia Stream water that originates in our upper watershed, then move through invasive wetland vegetation, areas undergoing invasive mangrove removal, and restored taro fields, and flow out towards a coastal ancient Hawaiian fishpond and out into Kāne'ohe Bay. The SWMP sites are intended to measure various parameters in real-time representing the emerging wetland/tidal marsh, estuary, coastal/ocean interface, and patch reef in the Bay. This indicates a gradient in salinity, temperature, and land-use change. We designed these SWMP sites to be able to track short- and long-term changes in our watershed through active restoration of habitat using a combination of conventional and Indigenous Native Hawaiian management strategies. As a measure of success, we hope to be able to capture the return to conditions ideal for many of our native species (fish, waterbirds, invertebrates, etc.) that have not been seen in the watershed for decades. We also envision the SWMP stations to serve as data hubs for researchers with various expertise to conduct question-specific research based around our consistent SWMP water quality measurements. Specifically, the SWMP nutrient dataset is designed to illustrate the nutrient consumption in the wetland and the fate of nutrients in the coastal fishpond and nearshore reef environments.

Our monthly grab sampling program consists of sampling for nutrients (nitrate + nitrite, orthophosphate, silicic acid, ammonium, total nitrogen and total phosphorus), total suspended solids, particulate carbon and nitrogen, chlorophyll *a*, and quarterly sampling of environmental DNA (eDNA) for fish.

For this 2022 submission, we report nutrient (nitrate + nitrite, orthophosphate, silicic acid, ammonia) and chlorophyll *a* data from four permanent SWMP stations (Wai 2, Reef 9, Kaho'okele, and Papahana).

b) Diel sampling program

We started our diel sampling program in November 2020 at Wai 2. Diel sampling was conducted at Wai 2 for November 2020 to January 2021. To accommodate an experiment looking at Chlorophyll sensors, diel sampling was moved to Kahoʻokele from February 2021 to April 2022. The diel sampling program returned to Wai 2 from May 2022 to current, to accommodate a grant project looking at tidal variability in the Heʻeia Stream.

3) Research methods -

a) Monthly grab sampling program

Sample collection and collection intervals: Before sampling, all nutrient bottles are rinsed 3 times with 10% hydrochloric acid and rinsed 3 times with DI water. Samples are collected on a monthly basis during the full moon phase on an outgoing tide (within 3 hours before slack low tide). Physical water parameters are documented using a YSI ProDSS immediately before the sample collection. Sampling is conducted by wading or free diving next to the deployed sonde and collecting water into two amber LDPE 500 mL bottles as close to the sensor depth as possible. Prior to collecting the sample water, each bottle is rinsed 3 times with ambient water, including with the cap. Time collected for each amber bottle is noted in our collection log sheet.

For Wai 2, samples collected before October 10, 2022 were obtained from the depth of the sensors, but at a location horizontally ~5 ft from the sonde, due to logistics of disturbing the site and impacting water quality parameters (specifically turbidity) when wading into the water to be near the sensor. However, since October 2022, we have decided to stand next to the sonde for ~3-5 min until the conditions settle down, pre-rinse the sample bottles, and open up the bottle at the depth of the sensors.

For Papahana, nutrients were collected monthly since 2018, prior to the SWMP station being deployed in June 2022. Prior to the station being established, grab samples were collected ~8-10 ft away, about a foot beneath the surface, in a freshwater pool that is expected to be homogeneous. Since the establishment of the sonde, nutrient samples have been collected immediately next to the sensors using the same method as described above. As we collect more data, we will determine if there are any statistical differences between samples collected prior to June 2022, and those collected currently.

Sample processing: Each amber 500 mL bottle is then processed for both nutrients and chlorophyll a. Using a syringe and a 25 mm GF/F filter in a filter holder, 5 mL of the filtrate is used to rinse the 60 mL LDPE preacid washed bottles twice prior to collecting 40 mL of the filtrate into the bottle for subsequent nutrient analyses. Then, another 50 mL is filtered through the GF/F filter for a total of 100 ml. The GF/F filter is then folded in half and stored in an aluminum foil packet. Both nutrients and chlorophyll samples, post collection, are then kept on ice in a cooler and delivered within 24 hours of field sampling to the University of Hawai'i SOEST Laboratory for Analytical Biogeochemistry (S-LAB), where they are frozen in a laboratory-grade walk-in -20°C freezer for subsequent laboratory processing. Subsequent analysese of nutrients are expected to occur within a month of receiving the samples.

Equipment and analyzers: Nutrients are processed on a Seal Analytical AA3 HR Nutrient Autoanalyzer following these established protocols. Chlorophyll *a* was analyzed on a Turner 10-AU fluorometer using the methods shown here and by Welschmeyer (1994). The 2023 WP-339 Final Report for the WatRTM Pollution Proficiency Testing Results (issued 06/05/2023) can be found here. The Method Detection Limits reported in this submission was analyzed using these EPA protocols. The Method Detection Limits used for each nutrient and chlorophyll *a* were calculated on June 12, 2023 and the results are reported here.

4) Site location and character –

He'eia (11.5 km²), O'ahu, Hawai'i, extends from the summit of the Ko'olau mountain range to the fourth largest wetland in the islands containing historically productive flooded taro agroecosystems (b'i), a coastal loko i'a, and into Kāne'ohe Bay highly valued for marine biodiversity. Ha'ikū and 'Ioleka'a basins contribute ~2.0 cfs perennial flow as He'eia Stream. State-wide, He'eia remains one of the few watersheds actively managed from ridge to reef. The He'eia NERR comprises a 1,385-acre region within the He'eia watershed, spanning the wetland and flooded-field agroecosystems managed by Kāko'o 'Ōiwi, the 800-year-old He'eia Fishpond managed by Paepae o He'eia, the surrounding streams and Kāne'ohe Bay under the jurisdiction of the state of Hawai'i Department of Land and Natural Resources, and Moku o Lo'e, the island on which Hawai'i Institute of Marine Biology, University of Hawai'i at Mānoa resides.

Wai 2 (W2): 21.43831° N, 157.81093° W

Wai 2, installed on September 18, 2019, is a site in He'eia Stream as the water flows out of the wetland area that has recently undergone large-scale mangrove removal. At W2, water either diverts and flows into the He'eia Fishpond, or goes straight towards the stream mouth into Kāne'ohe Bay. There is slight tidal influence, with intrusions of saline water at the peak of high tide when tide is greater than typical values. As a result, the typical salinity is 0.11 psu, but have observed salinity as high as 33.2 psu, very occasionally. Typical water depth is 0.52 m, increasing slightly only when heavy rains coincide with high tide. The bottom habitat is soft, silty, dark (anoxic) sediment. Pollutants in the area may include fecal contaminants from invasive cats, pigs, and mongoose, and potential cesspool contamination from the upper watershed. At W2, we now have an active data logger, and we are collecting monthly grab samples of nutrients, total suspended solids, chlorophyll *a*, and quarterly samples of eDNA.

Reef 9 (R9): 21.44628° N, 157.80183° W

Reef 9, installed on February 13, 2020, is a site on a patch reef located in Kāne'ohe Bay, a subtropical embayment on the windward coast of O'ahu. The salinity range is 33.7 psu to 35.0 psu, and depth changes ~1 m throughout tidal changes. This is a perpetual ocean site and presumed to have no input of freshwater except precipitation or during extreme low tide events. The bottom habitat is sandy, coral rubble, the average depth is ~3 m. Pollutants may include motorboat oil and human contaminants from nearby motorboat operators and proximity to He'eia Kea Small Boat Harbor.

Kaho'okele (KK): 21.43582° N, 157.80524W

Kahoʻokele, installed on September 29, 2020, is positioned on the ocean side (makai) of a ~800 year old Native Hawaiian fishpond. It is within close proximity to one of the fishpond sluice gates (mākāhā), also named Kahoʻokele, which serves as a point of water exchange between Heʻeia Stream and Kāneʻohe Bay. These traditional sluice gates are essential for regulating the physical-chemical parameters of the fishpond as well as maintenance of traditional husbandry of resource fish, and Kahoʻokele has been shown to exchange ~25% of the water exchanged through all the gates at the fishpond. The salinity range is 27.2 psu to 35.2 psu, and depth changes ~1 m throughout tidal changes. The bottom habitat is sandy, coral rubble, and pollutants may include pathogens related to domestic animal (i.e. cat, pig, mongoose) and other land-based pollutants coming from Heʻeia Stream and nearby Heʻeia State Park. Additionally, this site may be heavily influenced by groundwater sources of nutrients.

Papahana (PH): 21.40921° N, 157.82323° W

Papahana, installed on June 27, 2022, is located at Ha'akolea, or nicknamed Ice Pond, in Waipao in the back of the He'eia ahupua'a. Access is through the Indigenous organization Papahana Kuaola. The sensor, installed with telemetry, is co-located with a US Geological Survey stream gauge (16275000). It is a strictly freshwater site with no tidal influence. The bottom habitat is smooth river rock and soft sediment, with occasional flooding from seasonal rains. The water depth of the pool that the sonde sits in ranges from ~ 0.1 m typically, with much deeper conditions during heavy rains.

SWMP Station Timeline

Station Code	SWMP Status	Station Name	Location	Active Dates	Reason Decommissioned	Notes
W2	Р	Wai 2	21.43831° N, 157.81093° W	9/18/2019 - present	NA	NA
R9	Р	Reef 9	21.44628° N, 157.80183° W	2/13/2020 - present	NA	NA
KK	Р	Kahoʻokele	21.43582° N, 157.80524° W	9/29/2020- present	NA	NA

PH	P	Papahana	21.40921° N, 157.82323° W	6/27/2022- present	NA	NA
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5) Coded variable definitions –

heew2nut = He'eia Wai 2 Nutrients

heer9nut = He'eia Reef 9 Nutrients

heekknut = He'eia Kaho'okele Nutrients

heephnut = He'eia Papahana Nutrients

monthly grab sample program = 1 diel grab sample program = 2

6) Data collection period –

Wai 2 (W2):

Grab samples:

1/19/2022 10:13 10:14

2/16/2022 8:59 9:00

3/18/2022 8:47 8:49

4/14/2022 7:21 7:22

5/17/2022 8:07 8:08

6/15/2022 7:50 7:51

7/13/2022 7:35 7:36

8/10/2022 7:00 7:01

9/8/2022 6:37 6:38

10/10/2022 7:55 7:54

11/8/2022 8:04 8:05

12/7/2022 8:25 8:26

Diel samples:

5/15/2022 8:00 10:15 12:30, 14:45, 17:00, 19:15, 21:30, 23:45

5/16/2022 2:00, 4:15, 6:30, 8:45

6/14/2022 13:00, 15:15, 17:30, 19:45, 22:00

6/15/2022 0:15, 2:30, 4:45, 7:00, 9:15 (2 samples missing due to accidental discard while processing)

7/13/2022 8:45, 11:00, 13:15, 15:30, 17:45, 20:00 (6 samples missing due to ISCO battery malfunction, sampling redone for July on July 27-28)

7/27/2022 8:00, 10:15, 12:30, 14:45, 17:00, 19:15, 21:30, 23:45 7/28/2022 2:00, 4:15, 6:30, 8:45

8/10/2022 7:30, 9:45, 12:00, 14:15, 16:30, 18:45, 21:00, 23:15 8/11/2022 1:30, 3:45, 6:00, 8:15

9/8/2022 7:30, 9:45, 12:00, 14:15, 16:30,18:45, 21:00, 23:15 9/9/2022 1:30, 3:45, 6:00, 8:15

10/10/2022 9:30, 11:45, 14:00, 16:15, 18:30, 20:45, 23:00 10/11/2022 1:15, 3:30, 5:45, 8:00, 10:15

11/8/2022 9:45, 12:00, 14:15, 16:30, 18:45, 21:00, 23:15 11/9/2022 1:30, 3:45, 6:00, 8:15, 10:30

12/7/2022 3:30, 5:45, 8:00, 10:15, 12:30, 14:45, 17:00, 19:15, 21:30, 23:45 12/8/2022 2:00, 4:15

Reef 9 (R9):

Grab samples:

1/18/2022 9:00 9:01

2/15/2022 8:23 8:24

3/16/2022 7:37 7:38

4/13/2022 7:02 7:03

5/18/2022 8:48 8:49

6/14/2022 7:34 7:35

7/14/2022 7:52 7:53

8/11/2022 7:37 7:38

9/9/2022 7:04 7:05

10/11/2022 8:25 8:26

11/9/2022 8:34 8:35

12/8/2022 8:42 8:43

Kahoʻokele (KK):

Grab samples:

1/19/2022 9:37 9:38

2/16/2022 8:18 8:19

3/18/2022 7:49 7:50

4/14/2022 6:47 6:52

5/17/2022 7:21 7:22

6/15/2022 7:19 7:20

7/13/2022 6:48 6:49

8/10/2022 7:33 7:34

9/8/2022 6:49 6:50

10/10/2022 7:57 7:58

11/8/2022 8:10 8:11

12/7/2022 8:27 8:28

Diel samples:

1/18/2022 11:00,13:15, 15:30, 17:45, 20:00, 22:15

1/19/2022 0:30 2:45 5:00 7:15 9:30 11:45

2/15/2022 9:30, 11:45, 14:00, 16:15, 18:30, 20:45, 23:00

2/16/2022 1:15, 3:30, 5:45, 8:00 ,10:15

3/17/2022 9:15, 11:30, 13:45, 16:00, 18:15, 20:30, 22:45

3/18/2022 1:00, 3:15, 5:30, 7:45, 10:00

4/18/2022 10:15, 12:30, 14:45, 17:00, 19:15, 21:30, 23:45

4/19/2022 2:00, 4:15, 6:30, 8:45, 11:00

Papahana (PH):

Grab samples:

1/19/2022 12:44 12:45

2/16/2022 11:11 11:12

3/17/2022 11:19 11:20

4/14/2022 9:10 9:11

5/17/2022 11:17 11:18

6/15/2022 10:09 10:10 7/13/2022 9:23 9:24 8/10/2022 9:01 9:02 9/8/2022 8:29 8:30 10/10/2022 9:03 9:04 11/8/2022 9:28 8:30 12/7/2022 10:01 10:02

7) Associated researchers and projects-

As part of the SWMP long-term monitoring program, He'eia NERR also monitors 15-minute meteorological and water quality data which may be correlated with this nutrient/pigment dataset. These data are available at www.nerrsdata.org.

In addition to the SWMP stations, we also collect nutrients and pigments from 6 other stations in the He'eia watershed, designed to fill knowledge gaps on the fate of nutrients through the watershed. Many of these stations may be added as secondary SWMP stations in the future.

8) Distribution -

NOAA retains the right to analyze, synthesize and publish summaries of the NERRS System-wide Monitoring Program data. The NERRS retains the right to be fully credited for having collected and processed the data. Following academic courtesy standards, the NERR site where the data were collected should be contacted and fully acknowledged in any subsequent publications in which any part of the data are used. The data set enclosed within this package/transmission is only as good as the quality assurance and quality control procedures outlined by the enclosed metadata reporting statement. The user bears all responsibility for its subsequent use/misuse in any further analyses or comparisons. The Federal government does not assume liability to the Recipient or third persons, nor will the Federal government reimburse or indemnify the Recipient for its liability due to any losses resulting in any way from the use of this data.

Requested citation format:

NOAA National Estuarine Research Reserve System (NERRS). System-wide Monitoring Program. Data accessed from the NOAA NERRS Centralized Data Management Office website: www.nerrsdata.org; accessed 12 October 2022.

NERR nutrient data and metadata can be obtained from the Research Coordinator at the individual NERR site (please see Principal investigators and contact persons), from the Data Manager at the Centralized Data Management Office (please see personnel directory under the general information link on the CDMO home page) and online at the CDMO home page www.nerrsdata.org. Data are available in comma separated version format.

II. Physical Structure Descriptors

9) Entry verification –

Nutrient data is received by the Research Coordinator from the S-LAB Laboratory Specialist in a Microsoft Excel format. The original nutrient data is then archived in a shared Google Drive folder and an external hard drive. The S-LAB calculates and reports results in µg/L. For purposes of consistency in the NERR System, He'eia NERR converts the concentrations into mg/L by dividing all values by 1000. The SWMP Technician then enters the data from each month into the Microsoft Excel worksheet that will be subsequently processed

for QAQC. Prior to running NutrientQAQC, the worksheet is sent to the Research Coordinator for verification.

Nutrient data are entered into a Microsoft Excel worksheet and processed using the NutrientQAQC Excel macro. The NutrientQAQC macro sets up the data worksheet, metadata worksheets, and MDL worksheet; adds chosen parameters and facilitates data entry; allows the user to set the number of significant figures to be reported for each parameter and rounds using banker's rounding rules; allows the user to input MDL values and then automatically flags/codes measured values below MDL and inserts the MDL; calculates parameters chosen by the user and automatically flags/codes for component values below MDL, negative calculated values, and missing data; allows the user to apply QAQC flags and codes to the data; produces summary statistics; graphs selected parameters for review; and exports the resulting data file to the CDMO for tertiary QAQC and assimilation into the CDMO's authoritative online database.

10) Parameter titles and variable names by category -

Required NOAA NERRS System-wide Monitoring Program nutrient parameters are denoted by an asterisk "**"

Data Category	Parameter	Variable Name	Units of Measure
Phosphorus and	l Nitrogen:		
•	*Orthophosphate, Filtered	PO4F	mg/L as P
	*Ammonium, Filtered	NH4F	mg/L as N
	*Nitrite + Nitrate, Filtered	NO23F	mg/L as N
	Dissolved Inorganic Nitrogen	DIN	mg/L as N
Plant Pigments:			-
	*Chlorophyll a	CHLA_N	$\mu g/L$
Other Lab Para	meters:		
	Silicate, Filtered	SiO4F	mg/L as SI
Field Parameter	s:		
	Water Temperature	WTEM_N	°C
	Specific Conductance	SCON_N	mS/cm
	Salinity	SALT_N	ppt
	% Dissolved Oxygen Saturation	DO_S_N	0/0
	Dissolved Oxygen	DO_N	mg/L
	рН	PH_N	SU
	Turbidity – Nephelometer Turbidity Units	TURB_N	NTU

Notes:

- 1. Time is coded based on a 2400 clock and is referenced to Standard Time.
- 2. NO2 and NO3 are not reported individually and are reported together as NO23, as NO2 is a minor component (in the 0-1 nanomolar range) in Hawai'i's ocean and rain-fed systems relative to NO3 (https://hahana.soest.hawaii.edu/hot/hot-dogs/interface.html).

11) Measured or calculated laboratory parameters –

a) Parameters measured directly

Nitrogen species: NH4F, NO23F

Phosphorus species: PO4F

Other: CHLA_N, SiO4F

b) Calculated parameters

12) Limits of detection -

Method Detection Limits (MDL), the lowest concentration of a parameter that an analytical procedure can reliably detect, have been established by the S-LAB using these <u>EPA protocols</u>. The MDLs used for each nutrient and chlorophyll *a* were calculated June 2023 and the results are reported <u>here</u>. As the results were reported in μg/L, they were divided by 1000 to yield concentrations in mg/L as NO23, NH4, PO4, and SiO4. Different MDLs were reported for 2 separate (lower and higher) concentration ranges for NO23, PO4, and SiO4, and we selected the value that was most appropriate for our specific dataset. The CHLA MDL was calculated as 3 times the standard deviation of the blank (n=15), documentation can be found <u>here</u>. These values are reviewed and revised annually.

Parameter	Start Date	End Date	MDL	Revisited
NH4F	01/01/22	12/31/22	0.00042	
NO23F	01/01/22	12/31/22	0.0001	
PO4F	01/01/22	12/31/22	0.00028	
SiO4F	01/01/22	12/31/22	0.00225	
CHLA_N	01/01/22	12/31/22	0.023	

13) Laboratory methods –

All dissolved nutrient samples are stored frozen until analysis. Samples are thawed overnight in a refrigerator. Standards are made from concentrated stock solutions daily. For determination of NO23F and TN, a NO2 and NO3 is included in the run to determine efficiency of cadmium column reduction. For TN and TP determination an organic TN and TP sample is included in the run to determine efficiency of UV digestion. Every 15 samples the following sequence is completed: deionized baseline, deionized water blank, low nutrient seawater blank, two replicate standards at the middle concentration (to determine consistent recovery and accuracy), a low concentration QA/QC sample, a high concentration QA/QC sample, and finally, the high standard (to determine instrument drift).

QA/QC samples analyzed throughout the run are collected by the <u>Hawaii Ocean Time-series program</u>. These are collected with the CTD-Niskin bottles at 250 m and 4000 m. The 250 m sample acts as the high concentration sample for total dissolved nitrogen and total dissolved phosphorus and acts as the low concentration sample for the suite of inorganic dissolved nutrients. The 4000 m sample acts as the low concentration sample for total dissolved nitrogen and total dissolved phosphorus and acts as the high concentration sample for the suite of inorganic dissolved nutrients.

Refer to document "S-LAB Description of Procedures" on file with CDMO for specific procedures and references.

a) Parameter: NH4F

S-LAB Laboratory Method: S-LAB Description of Procedures

Method Reference: Kerouel and Aminot (1997)

Method Descriptor: The sample is reacted with o-phthaladehyde (OPA) at 75°C in the presence of borate buffer and sodium sulfite to form a fluorescent species proportional to the ammonia concentration. The fluorescence is measured at 460 nm following excitation at 370 nm.

Preservation Method: Samples filtered and stored at -20 °C up to 28 days.

b) Parameter: NO23F

S-LAB Laboratory Method: S-LAB Description of Procedures
Method Reference: Armstrong et al. (1967); Grasshof (1969); Grasshof et al. (1983)

Method Descriptor: The determination of nitrate and nitrite uses a procedure whereby nitrate is reduced to nitrite by a copper-cadmium redactor column. The nitrite then reacts with sulfanilamide under acidic conditions to form a diazo compound. This compound then couples with N-1-napthylethylene diamine dihydrochloride to form a purple azo dye. The method is based on the nitrate determination in Standard Methods and in the DIN/ISO Standards for automatic nitrate measurements. Detection is at 520 nm.

Preservation Method: Samples filtered and stored frozen at -20 °C up to 28 days.

c) Parameter: PO4F

S-LAB Laboratory Method: S-LAB Description of Procedures

Method Reference: Grasshoff (1965)

Method Descriptor: Automated procedure for the determination of orthophosphate based on the colorimetric method in which blue color is formed by the reaction orthophosphate, molybdate ion and antimony ion followed by reduction with ascorbic acid at a pH<1. The reduced blue phosphor-molybdenum complex is

colorimetrically read at 880 nm.

Preservation Method: Samples filtered and stored at -20 °C up to 28 days.

d) Parameter: SiO4F

S-LAB Laboratory Method: S-LAB Description of Procedures

Method Reference: Grasshoff et al. (1983)

Method Descriptor: This automated procedure for the determination of soluble silicates is based on the reduction of silicomolybdate in acidic solution to molybdenum blue by ascorbic acid. Oxalic acid is introduced to the sample stream before the addition of ascorbic acid to minimize interference from phosphates. Detection is at 820 nm.

Preservation Method: Samples filtered and stored at -20 °C up to 28 days.

e) Parameter: CHLA_N

S-LAB Laboratory Method: S-LAB Description of Procedures

Method Reference: Welschmeyer (1994)

Method Descriptor: Samples are extracted with a 90% acetone solution at the end of the day (after 3 PM). They are stored in the freezer overnight for extraction. Sample extraction ranges between 16-19 hours. Sample extraction is never less than 2 hours or greater than 24 hours. Samples are brought to room temperature prior to analysis. At the start and end of each day a high and low solid standard is analyzed on the Turner 10AU fluorometer to monitor any long-term instrument drift. The Turner 10 AU fluorometer is calibrated every 6 months or as needed based on solid standard drift (not to exceed 5%). A blank of 90% acetone is analyzed at the start and end of each day. Samples are homogenized and the 90% acetone solution is decanted into a new borosilicate test tube for analysis. If dilution is required, a known volume of the original sample is pipetted into a new test tube, diluted with a known volume of 90% acetone, homogenized, and analyzed. The Turner 10 AU is equipped with a non-acidification module that does not require acidification to account for chlorophyll b and pheopigments

Preservation Method: Samples filtered and stored at -20 °C up to 30 days.

14) Field and Laboratory QAQC programs –

a) Precision

i) Field variability -

Field replicates:

<u>Nutrients</u>: 46 replicates / 94 total samples =49% <u>Chlorophyll:</u> 46 replicates / 94 total samples =49%

<u>For both nutrients and chlorophyll, true</u> field replicates were collected for all sites for all months in 2023, by collecting successive grab samples (separate sample in another sample bottle from the field at a different time than the first replicate).

				Nutrien	ts		Chlorop	hyll
Site	Date	Collection times	Sample	Split	Replicate	Sample	Split	Replicate
Wai 2	1/19/2022	10:13 10:14	1		1	1		1
	2/16/2022	8:59 9:00	1		1	1		1
	3/18/2022	8:47 8:49	1		1	1		1
	4/14/2022	7:21 7:22	1		1	1		1
	5/17/2022	8:07 8:08	1		1	1		1
	6/15/2022	7:50 7:51	1		1	1		1
	7/13/2022	7:35 7:36	1		1	1		1
	8/10/2022	7:00 7:01	1		1	1		1
	9/8/2022	6:37 6:38	1		1	1		1
	10/10/2022	7:55 7:54	1		1	1		1
	11/8/2022	8:04 8:05	1		1	1		1
	12/7/2022	8:25 8:26	1		1	1		1

				Nutrien	ts		Chloroph	yll
Site	Date	Collection times	Sample	Split	Replicate	Sample	Split	Replicate
Reef 9	1/18/2022	9:00 9:01	1			1		
	2/15/2022	8:23 8:24	1			1		
	3/16/2022	7:37 7:38	1		1	1		1
	4/13/2022	7:02 7:03	1		1	1		1
	5/18/2022	8:48 8:49	1		1	1		1
	6/14/2022	7:34 7:35	1		1	1		1
	7/14/2022	7:52 7:53	1		1	1		1
	8/11/2022	7:37 7:38	1		1	1		1
	9/9/2022	7:04 7:05	1		1	1		1
	10/11/2022	8:25 8:26	1		1	1		1
	11/9/2022	8:34 8:35	1		1	1		1
	12/8/2022	8:42 8:43	1		1	1		1

				Nutrients			Chloroph	yll
Site	Date	Collection times	Sample	Split	Replicate	Sample	Split	Replicate
Kahoʻokele	1/19/2022	9:37 9:38	1		1	1		1
	2/16/2022	8:18 8:19	1		1	1		1
	3/18/2022	7:49 7:50	1		1	1		1

4/14/2022	6:47 6:52	1	1	1	1
5/17/2022	7:21 7:22	1	1	1	1
6/15/2022	7:19 7:20	1	1	1	1
7/13/2022	6:48 6:49	1	1	1	1
8/10/2022	7:33 7:34	1	1	1	1
9/8/2022	6:49 6:50	1	1	1	1
10/10/2022	7:57 7:58	1	1	1	1
11/8/2022	8:10 8:11	1	1	1	1
12/7/2022	8:27 8:28	1	1	1	1

				Nutrien	ts	(Chloroph	yll
Site	Date	Collection times	Sample	Split	Replicate	Sample	Split	Replicate
Papahana	1/19/2022	12:44 12:45	1		1	1		1
	2/16/2022	11:11 11:12	1		1	1		1
	3/17/2022	11:19 11:20	1		1	1		1
	4/14/2022	9:10 9:11	1		1	1		1
	5/17/2022	11:17 11:18	1		1	1		1
	6/15/2022	10:09 10:10	1		1	1		1
	7/13/2022	9:23 9:24	1		1	1		1
	8/10/2022	9:01 9:02	1		1	1		1
	9/8/2022	8:29 8:30	1		1	1		1
	10/10/2022	9:03 9:04	1		1	1		1
	11/8/2022	9:28 8:30	1		1	1		1
	12/7/2022	10:01 10:02	1		1	1		1

- ii) Laboratory variability 0 Laboratory replicates / 94 total samples = 0%
- iii) Inter-organizational splits Samples were not split and analyzed by two different labs.

b) Accuracy

- i) Sample spikes Standards utilized are "spiked" low nutrient seawater (LNSW), though no samples are spiked. Middle LNSW spiked standards are evaluated in the same way, the % recovery of this is reported every 15 samples (duplicate Cal 3 on each run file), and % recovery is typically 100% +/- 5%.
- ii) **Standard reference material analysis** Certified reference materials (NMIJ Seawater Standards) are analyzed during each nutrient run, and the S-LAB values are consistently within the accepted ranges. The 2023 ERA's WP-339 WatRTMPollution Proficiency Testing using standards was conducted April to June 2023 is found https://example.com/here.com/h
- iii) Cross calibration exercises –He'eia NERR has not participated in any cross calibration exercises.

15) QAQC flag definitions –

QAQC flags provide documentation of the data and are applied to individual data points by insertion into the parameter's associated flag column (header preceded by an F_). QAQC flags are applied to the nutrient data during secondary QAQC to indicate data that are out of sensor range low (-4), rejected due to QAQC checks (-3), missing (-2), optional and were not collected (-1), suspect (1), and that have been corrected (5). All remaining data are flagged as having passed initial QAQC checks (0) when the data are uploaded and assimilated into the CDMO ODIS as provisional plus data. The historical data flag (4) is used to indicate data that were submitted to the CDMO prior to the initiation of secondary QAQC flags and codes (and the use of the automated primary QAQC system for WQ and MET data). This flag is only present in historical data that are exported from the CDMO ODIS.

- -4 Outside Low Sensor Range
- -3 Data Rejected due to QAQC
- -2 Missing Data
- -1 Optional SWMP Supported Parameter
- 0 Data Passed Initial QAQC Checks
- 1 Suspect Data
- 4 Historical Data: Pre-Auto QAQC
- 5 Corrected Data

16) QAQC code definitions -

QAQC codes are used in conjunction with QAQC flags to provide further documentation of the data and are also applied by insertion into the associated flag column. There are three (3) different code categories, general, sensor, and comment. General errors document general problems with the sample or sample collection, sensor errors document common sensor or parameter specific problems, and comment codes are used to further document conditions or a problem with the data. Only one general or sensor error and one comment code can be applied to a particular data point. However, a record flag column (F_Record) in the nutrient data allows multiple comment codes to be applied to the entire data record.

General errors

GCM	Calculated value could not be determined due to missing data
GCR	Calculated value could not be determined due to rejected data
GDM	Data missing or sample never collected
GQD	Data rejected due to QA/QC checks
GQS	Data suspect due to QA/QC checks
GSM	See metadata

Sensor errors

SBL	Value below minimum limit of method detection
SCB	Calculated value could not be determined due to a below MDL component
SCC	Calculation with this component resulted in a negative value
SNV	Calculated value is negative
SRD	Replicate values differ substantially
SUL	Value above upper limit of method detection

Parameter Comments

CAB	Algal bloom
CDR	Sample diluted and rerun
CHB	Sample held beyond specified holding time
CIP	Ice present in sample vicinity
CIF	Flotsam present in sample vicinity
CLE	Sample collected later/earlier than scheduled

```
CRE Significant rain event
```

CSM See metadata

CUS Lab analysis from unpreserved sample

Record comments

- CAB Algal bloom
- CHB Sample held beyond specified holding time
- CIP Ice present in sample vicinity
 CIF Flotsam present in sample vicinity
- CLE Sample collected later/earlier than scheduled
- CRE Significant rain event
- CSM See metadata
- CUS Lab analysis from unpreserved sample

Cloud cover

- CCL clear (0-10%)
- CSP scattered to partly cloudy (10-50%)
- CPB partly to broken (50-90%)
- COC overcast (>90%)
- CFY foggy
- CHY hazy
- CCC cloud (no percentage)

Precipitation

- PNP none PDR drizzle PLR light rain
- PHR heavy rain
 PSQ squally
- PFQ frozen precipitation (sleet/snow/freezing rain)
- PSR mixed rain and snow

Tide stage

- TSE ebb tide TSF flood tide
- TSH high tide
- TSL low tide

Wave height

- WH0 0 to <0.1 meters
- WH1 0.1 to 0.3 meters
- WH2 0.3 to 0.6 meters
- WH3 0.6 to > 1.0 meters
- WH4 1.0 to 1.3 meters
- WH5 1.3 or greater meters

Wind direction

- N from the north
- NNE from the north northeast
- NE from the northeast
- ENE from the east northeast
- E from the east
- ESE from the east southeast SE from the southeast
- SSE from the south southeast
- S from the south
- SSW from the south southwest
- SW from the southwest
- WSW from the west southwest

W from the west
WNW from the west northwest
NW from the northwest
NNW from the north northwest
Wind speed
WS0 0 to 1 knot
WS1 > 1 to 10 knots

WS1 > 1 to 10 knots

WS2 > 10 to 20 knots

WS3 > 20 to 30 knots

WS4 > 30 to 40 knots

WS5 > 40 knots

17) Other remarks/notes -

Data may be missing due to problems with sample collection or processing. Laboratories in the NERRS System submit data that are censored at a lower detection rate limit, called the Method Detection Limit or MDL. MDLs for specific parameters are listed in the Laboratory Methods and Detection Limits Section (Section II, Part 12) of this document. Concentrations that are less than this limit are censored with the use of a QAQC flag and code, and the reported value is the method detection limit itself rather than a measured value. For example, if the measured concentration of NO23F was 0.0005 mg/l as N (MDL=0.0008), the reported value would be 0.0008 and would be flagged as out of sensor range low (-4) and coded SBL. In addition, if any of the components used to calculate a variable are below the MDL, the calculated variable is removed and flagged/coded -4 SCB. If a calculated value is negative, it is rejected and all measured components are marked suspect. If additional information on MDL's or missing, suspect, or rejected data is needed, contact the Research Coordinator at the reserve submitting the data.

Note: The way below MDL values are handled in the NERRS SWMP dataset was changed in November of 2011. Previously, below MDL data from 2007-2010 were also flagged/coded, but either reported as the measured value or a blank cell. Any 2007-2011 nutrient/pigment data downloaded from the CDMO prior to November of 2011 will reflect this difference.

Sample hold times for 2022: Samples are held at -20°C. NERRS SOP allows nutrient samples to be held for up to 28 days (CHLA for 30) at -20°C, plus allows for up to 5 days for collecting, processing, and shipping samples. Samples held beyond that time period are flagged suspect <1>and coded (CHB). If measured values were below MDL, this resulted in <-4> [SBL] (CHB) flagging/coding.

*Sample held longer than allowed by NERRS protocols

	Date of analysis					
Sample Descriptor	PO4F	NH4F	NO23F	CHLA_N	SiO4F	
1/18, 1/19/2022, all grab and diel samples	1/31/2022	1/31/2022	1/31/2022	1/26/2022	1/31/2022	
2/15, 2/16/2022, all grab and diel samples	2/17/2022	2/17/2022	2/17/2022	2/22/2022 (grab) 2/23/2022 (diel)	2/17/2022	
03/17, 3/18/2022, all grab and diel samples	3/21/2022	3/21/2022	3/21/2022	3/29/2022	3/21/2022 3/22/2022	
4/13, 4/14/2022, all grab samples	*5/19/2022	*5/19/2022	*5/19/2022	*5/18/2022	*5/19/2022	

4/13, 4/14/2022, all diel	*6/27/2023	*6/27/2023	*6/27/2023	6/13/2023	*6/27/2023
samples					
5/17, 5/18/2022 all grab	*6/27/2022	*6/27/2022	*6/27/2022	6/13/2022	*6/27/2022
and diel samples					
6/14, 6/15/2022, all grab	7/08/2022	7/08/2022	7/08/2022	7/01/2022	7/08/2022
and diel samples					
7/13, 7/14/2022, all grab	*8/15/2022	*8/15/2022	*8/15/2022	8/09/2022	*8/15/2022
and diel samples					
8/10, 8/11/2022, all grab	8/30/2022	8/30/2022	8/30/2022	*9/14/2022	8/30/2022
and diel samples				(grab)	
				8/18/2022	
				(diel)	
9/08, 9/09/2022, all grab	9/20/2022	9/20/2022	9/20/2022	9/14/2022	9/20/2022
and diel samples					
10/10, 10/11/2022, all	10/12/2022	10/12/2022	10/12/2022	10/25/2022	10/12/2022
grab and diel samples				(grab)	
				10/24/2022	
				(diel)	
11/08, 11/09/2022, all	*12/12/2022	*12/12/2022	*12/12/2022	12/01/2022	*12/12/2022
grab and diel samples					
12/07, 12/08/2022, all	12/20/2022	12/20/2022	12/20/2022	1/04/2023	12/20/2022
grab and diel samples				(grab)	
				1/05/2023	
				(diel)	

Site

Date Code 1/19/2022 CAB

Note

Significantly low silicate concentrations observed in our marine stations R9 and KK, associated with relatively higher CHL concentrations for both R9 and KK, indicative of an algae (possibly diatom) bloom. While the timeline does not quite match up, this "bloom" could have been due to landward influence of a large rain event (>400 cfs stream flow at Papahana) that occurred 2 weeks prior, on Dec 30, 2021.

1/19/2022 CSM

Prior to the installation of the Papahana station, sampling was conducted ~7-8 ft away from the SWMP station, but in the same pool. Following the SWMP station installation on 06/27/22, sampling was conducted immediately adjacent to the sensors on the sonde, in a notably deeper section of the stream/pool. While this change in the precise sample location did not seem to change the typical nutrient concentrations or physical parameters of the water body, we will keep conduct statistical analyses once we acquire more data. Historical nutrient data (from 2018) for this site will be submitted and checked to account for seasonal variability.

2/16/2022	CSM	Same note as above
3/17/2022	CSM	Same note as above
4/14/2022	CSM	Same note as above
5/17/2022	CSM	Same note as above
6/15/2022	CSM	Same note as above
8/11/2022 7:38	-2 CSM	A missing chl sample at reef 9 that may have gotten lost at the S-lab analytical lab processing stage.