Sapelo Island (SAP) NERR Nutrient Metadata January 2008 - December 2008 Latest Update: June 12, 2015

I. Data Set and Research Descriptors

1) Principal investigator(s) and contact persons

a) Reserve Contact

Dorset Hurley P.O. Box 15 Sapelo Island, GA 31327

Sapelo Island, GA 31327 Phone: 912-485-2251

e-mail: dhurley@darientel.net

b) Laboratory Contact

Katy Austin Smith
715 Bay Street
Marine Extension Service Laboratory
University of Georgia
Brunswick, GA 31520

Phone: 912-262-3338 e-mail: klaustin@uga.edu

c) Other Contacts and Programs

none

2) Research objectives

The nutrient monitoring program is designed upon spatial deployment across a wide variety of marsh types with differing fresh and marine water mixing. These differing dynamics allow scientists and researchers to select from both a wide variety of research sites as well as tailor research programs to specific tidal dynamics and utilize the Reserves SWMP data acquisitions to the maximum extent. Additionally, from a long-term trend perspective the variety of marsh types and hydrology being monitored will allow for a better understanding of the different effects of sea-level rise upon marsh type. Due to a lack of residential development and very low human activity within the watersheds of the sites, they serve as a proxy for reference conditions with the various marsh and associated hydrology types for the creeks and river stations. All of the sites selected have very little anthropogenic nutrient influences. The following brief descriptions are associated with each nutrient monitoring site. For more detail please refer to the site descriptors located under section (4) of this document and/or contact the Research Coordinator at the SAP NERR for detailed information of any/all sites.

<u>Lower Duplin</u>: Located at the mouth of the Duplin River with large, rapid and near-complete hydraulic exchange with Doboy Sound within each diurnal cycle. Typical of a high salinity, well mixed estuary site.

<u>Hunt Dock</u>: Located on the upper Duplin with relatively high hydraulic retention requiring an estimated 6-7 diurnal events to complete a total hydraulic exchange. Rainfall may drop salinity precipitously in the basin depending on tidal height, duration and volume of precipitation.

<u>Cabretta Creek</u>: Located on the eastern side of Sapelo Island with direct exchange with the Atlantic Ocean. Creek is typical of high salinity, high oceanic exchange and near complete hydraulic exchange with each diurnal event. Creek is extremely buffered from rainfall (event driven) fluctuations in salinity.

<u>Dean Creek:</u> Located on the southern end of Sapelo is the primary drainage of the inter-dune (ocated amid primary and secondary dune systems) meadow. This site is highly susceptible to very high salinity fluctuations associated with rainfall events on both seasonal and short —term, event driven scales. Tidal exchange is complete at each diurnal event and exchange water genesis is the Doboy Sound.

The Duplin River is a tidal basin with no freshwater influence within its headwaters apart from surficial aquifer weeping from the perched lens of water associated with Sapelo Island. This nutrient monitoring effort is tied into the Georgia Coastal Ecosystems, Long-Term Ecological Research (GCE-LTER) initiative and the University of Georgia Marine Extension Service water quality database whose collection and analysis of the water samples facilitates the database. This long-term data set is being developed to provide information on estuarine water mixing within the well-studied Duplin River basin in addition to providing a long-term characterization of water quality as related to nutrient loading within the Duplin River.

- a) **Monthly Grab** The monthly grab sampling program focuses on documentation of baseline reference nutrient trends within a wide array of local marsh systems with differing hydrology.
- b) **Diel Sampling Program** The diel sampling program focuses on short-term temporal variability over a lunar tidal cycle.

3) Research methods

a) Monthly Grab Sampling Program

Monthly grab samples were taken at four stations within the Duplin River estuary from January to December 2008. Bottom water samples were taken at the Lower Duplin (LD), Hunt Dock (HD), Cabretta Creek (CA) and Dean Creek (DC) stations using a Niskin style sampling bottle. All grab samples were taken sequentially in duplicate beginning near the time the last diel sample was collected by the ISCO sampler (this time corresponds to low tide at the end of the tidal cycle). Chronological collection times for each of the four sites varied as two teams of people were conducting the actual sampling. A group of two remained onboard the research vessel to sample at the Hunt Dock and Lower Duplin sites, while the other group drove inland to Cabretta Creek and Dean Creek. At the time of sample collection, latitude, longitude, time and depth were recorded. All grab samples were collected from the Niskin bottle into an acid-washed (10% HCl) polypropylene beaker for filtering. Two filter towers were set up, one acid-washed tower with a 0.45 um polycarbonate filter for nutrient filtering and one clean tower with a GF/F filter for chlorophyll filtering. A small amount of sample was used to rinse the nutrient filter tower equipped with a filter and then the filtrate was discarded. The tower was then filled to the 250mL mark. The chlorophyll tower with the GF/F filter was also filled to the 250-mL mark and the two towers were connected by a small piece of tubing. The vacuum pump was turned on to pull

the 250 mL through each filter and then the vacuum was released. The nutrient sample tower was disconnected and an acid-washed 250-mL polypropylene bottle was rinsed and filled with the filtrate. Space was left in the sample bottle for expansion during freezing at approximately -18 degC. If the first 250 milliliters went through the chlorophyll filter easily, the filtrate was discarded and an additional 50, 100 or 250 milliliters was filtered, depending on suspended sediment load, to concentrate the sample onto the filter. The chlorophyll filter was then removed with tweezers and placed face up in a petri dish, wrapped in aluminum foil and labeled with the volume filtered and sample information. The chlorophyll filter towers were rinsed between replicate grabs with distilled water and the nutrient filter tower was acid-washed and DI water rinsed between samples. Nutrient and chlorophyll filtering between grabs took approximately 10 minutes to complete. At the Cabretta and Dean Creek sites, a vacuum hand pump was used rather than a mechanical pump, which is available only on the research vessel. The depths at these two sites were estimated as sampling took place from a bridge. Samples were immediately placed on ice, in the dark and returned to the laboratory within six hours. Once in the laboratory, samples were frozen and processed within the specified times (unless flagged) for nutrient and chlorophyll-a concentrations.

b) Diel Sampling Program

WWW Tide and Current Predictor for Wolf Island, South End was used to estimate low tide. As close to an early, low, neap tide as possible was selected each month for sampling. The ISCO sampler was deployed at the Lower Duplin (LD) site on the day previous to the grab sample date chosen for that particular month with the sample line suction placed at 1.5 feet below the surface of the water. The ISCO sampler collected the first diel sample two hours later than low tide on the following day and continued collecting samples every two hours for the next 22 hours, representing a full tidal cycle, a total of 12 samples, and ending at low tide when grab sampling began. The ISCO was turned off and the 12 samples were secured with caps upon arriving at the site. The samples were filter processed either in the field after completion of grab sampling at Lower Duplin or back in the laboratory, weather depending. The filtration process for the diel samples follows the same process as for grab samples described above. High-density polypropylene bottles were used to collect the samples after filtration. Polypropylene bottles and filter towers were soaked in 10% HCl in preparation for the fieldwork, then triple rinsed with distilled water. A squeeze bottle was used to acid wash (then rinse with distilled water) beakers and filter towers in the field between filtering of each sample.

4) Site location and character

The Sapelo Island National Estuarine Research Reserve is located on the Southeastern Atlantic coast of the United States in McIntosh County, Georgia. The study area encompasses the Duplin River estuary, a tidally flushed drainage system flowing into Doboy Sound from the north and two inland creeks, Cabretta and Dean's Creek. The Duplin River watershed occupies most of the Reserve, which also contains various forest types, sand dunes, a section of ocean beach and minor developed areas. The Duplin River estuary covers 3,300 acres between Sapelo Island and the mainland in McIntosh County. It drains a tidal bay and an extensive network of salt marshes about 6 miles long, into which there is little upland run-off. Diverse estuarine wetlands provide extensive and complex habitat types for fish and wildlife. The island contains several small, interior brackish and freshwater marshes fed by surficial aquifer expression (interdune meadow of Nannygoat beach: south end) and anthropogenic upland ditches and dikes produced in the early 19th century (north end). The upland forests are composed of several diverse habitats including long leaf pine/slash pine forests, climax maritime forests, small amounts of pond cypress bays and naturally regenerated loblolly pine forests which are

timbered on a 70 year selectively cut harvest rotation. There are no current studies on pollutants in this area. Sapelo Island is typically considered a pristine environment, with minimal pollutant input.

Latitude and Longitude-

Lower Duplin: Lat: 31 25' 4" N, Long: 81 17' 46" W Hunt Dock: Lat: 31 28' 43" N, Long: 81 16' 23" W Cabretta Creek: Lat: 31 26 37.3" N, Long: 81 14 23.7" W Dean Creek: Lat: 31 23 22.5" N, Long: 81 16 44.2" W

Water Quality site descriptions-

Salinities at all Duplin River sites vary according to localized rainfall and associated runoff. The upper Duplin River site (Hunt Dock) experiences slightly lower salinities associated with rainfall events (2 -3ppt) as compared to the lower Duplin River site. Average salinities range from 15 ppt to 30 ppt depending on seasonal or event rainfall. Average tidal range of diurnal tidal cycle is approximately 2.5 meters twice daily. Due to high turbidity, all Duplin River sites are lacking any persistent submerged aquatic vegetation and have an unconsolidated sandy/mud bottom (soft sediment) typical of southeastern near-ocean estuaries. Marsh sediments are relatively pristine and free of pollutants based on sediment analysis conducted in 1996 by C. Alexander, Skidaway Institue of Oceanography. Watershed is dominated by oceanic tidal influences associated with Doboy Sound. Depths are as follows: Lower Duplin (LD) ranges from 1.5 meters to 6.0 meters depending on tide, and Hunt Dock's maximum depth is 4.27 meters.

Cabretta Creek is fed directly from waters of the Atlantic Ocean. Cabretta experiences a maximum tidal range of approximately 4.3 meters. Average mean low water depth at the sample site is approximately 3.25 meters. Salinity ranges, with exception to major, long-term precipitation events, from 15-36 ppt, seasonally. The station is located on a small (one-lane), wooden, roadway bridge spanning Cabretta Creek, located on the island's extreme eastern side. The benthos is composed primarily of sand substrate with small, intertidal oyster reef conglomerate communities. Adjacent to the site is extensive, intertidal, bank stabilization (armoring) in the form of woven rip-rap fencing and granite rocks. This manipulation is slowly becoming stabilized via oyster reef community colonization. The adjacent marshes are dominated by Spartina alterniflora with occasional Juncus romerianus in the nearby fringe community habitat. The creek has very little adjacent uplands due to: 1) the low elevational gradient and 2) the area's geologically recent accretion genesis (Holocene) resulting in sandy soils; of which neither condition allows for extensive floral colonization or stabilization.

The Dean Creek site is located on a small wooden bridge spanning Dean Creek, in close proximity to the adjacent Nannygoat Beach causeway. Dean Creek is a small tidal basin fed from the waters of Doboy Sound, which is located on Sapelo Island's south end. With exception to short duration local or long duration regional precipitation events, the creek's salinity normally ranges between 20 and 30 ppt. The benthic community consists of a sandy-mud substrate with occasional small, intertidal oyster reef community and mean tidal amplitude of approximately 8 feet. Average mean low water depth at the sample site is approximately 1 meter, but fluctuates due to bank erosion. The small creek feeds approximately 150 acres of Spartina alterniflora dominated salt marsh, which is interspersed with small 0.5-1 acre hammocks and salt pans. Fringe community components range from Loblolly pine forests with a sub-canopy of Yaupon holly to Wax myrtle and Sable Palm.

5) Coded variable definitions

LD = Lower Duplin; HD = Hunt Dock; CA = Cabretta Creek; DC = Dean Creek.

Each individual sample is given a 3 part name code in addition to other codes. The 3 part name code, "sapldnut" for example, gives the reserve name (sap = Sapelo), station name (LD = Lower Duplin, etc), and SWMP program code (nut = nutrient monitoring program).

Sampling Site codes:

sapldnut – Sapelo Island nutrient data for Lower Duplin saphdnut – Sapelo Island nutrient data for Hunt Dock sapcanut – Sapelo Island nutrient data for Cabretta Creek sapdcnut – Sapelo Island nutrient data for Dean Creek

The monitoring codes are set as "1" to indicate grab samples and "2" to indicate diel samples. Replicates are also given specific codes. Grab samples in which duplicate field samples are taken utilize a "1" for the first sample and a "2" for the second sample. Subsequent lab splits of each field rep are labeled with an "S". Diel samples are always labeled with a "1" for the first lab replicate and an "S" for the second lab replicate. Only one actual sample is taken at each interval with the ISCO sampler.

6) Data collection period

Diel sampling for 2008 began at 11:30:00 on January 28, 2008. Grab sampling commenced on January 29, 2008 at 09:40:00 at the Lower Duplin site. Due to inclement weather and subsequent instrument (ISCO sampler) malfunction, diel samples were not collected during February 2008.

Diel S	<u>Sampling</u>			
Site	Start	Start	End	End
	Date	Time	Date	Time
LD	01/28/2008	1130	01/29/2008	0930
LD	03/18/2008	1623	03/19/2008	1423
LD	04/15/2008	1507	04/16/2008	1307
LD	05/13/2008	1343	05/14/2008	1143
LD	06/10/2008	1210	06/11/2008	1010
LD	07/08/2008	1035	07/09/2008	0835
LD	08/11/2008	1406	08/12/2008	1206
LD	09/08/2008	1230	09/09/2008	1030
LD	10/06/2008	1045	10/07/2008	0845
LD	11/10/2008	1505	11/11/2008	1305
LD	12/08/2008	1345	12/09/2008	1145
Grab :	Sampling			
Site	Start	Start	End	End
	Date	Time	Date	Time
CA	01/29/2008	1006	01/29/2008	1017
CA	02/21/2008	1535	02/21/2008	1545
CA	03/19/2008	1437	03/19/2008	1449
CA	04/16/2008	1443	04/16/2008	1454

CA	05/14/2008	1311	05/14/2008	1324
CA	06/11/2008	1050	06/11/2008	1058
CA	07/09/2008	0928	07/09/2008	0940
		1224		1238
CA	08/12/2008		08/12/2008	
CA	09/09/2008	1115	09/09/2008	1129
CA	10/07/2008	1048	10/07/2008	1101
CA	11/11/2008	1401	11/11/2008	1414
CA	12/09/2008	1245	12/09/2008	1259
LD	01/29/2008	0940	01/29/2008	0952
LD	02/21/2008	1513	02/21/2008	1522
LD	03/19/2008	1357	03/19/2008	1408
LD	04/16/2008	1324	04/16/2008	1332
LD	05/14/2008	1200	05/14/2008	1210
LD	06/11/2008	0959	06/11/2008	1011
LD	07/09/2008	0956	07/09/2008	1002
LD	08/12/2008	1200	08/12/2008	1213
LD	09/09/2008	1047	09/09/2008	1054
LD	10/07/2008	1015	10/07/2008	1029
LD	11/11/2008	1338	11/11/2008	1352
LD	12/09/2008	1218	12/09/2008	1235
HD	01/29/2008	1016	01/29/2008	1025
HD	02/21/2008	1435	02/21/2008	1441
HD	03/19/2008	1316	03/19/2008	1325
HD	04/16/2008	1247	04/16/2008	1257
HD	05/14/2008	1120	05/14/2008	1126
HD	06/11/2008	0925	06/11/2008	0933
HD	07/09/2008	0915	07/09/2008	0922
HD	08/12/2008	1044	08/12/2008	1105
HD	09/09/2008	1010	09/09/2008	1018
HD	10/07/2008	0930	10/07/2008	0938
HD	11/11/2008	1233	11/11/2008	1244
HD	12/09/2008	1135	12/09/2008	1143
	12.03.2000	1100	12.09.2000	11.0
DC	01/29/2008	1050	01/29/2008	1100
DC	02/21/2008	1623	02/21/2008	1630
DC	03/19/2008	1524	03/19/2008	1533
DC	04/16/2008	1537	04/16/2008	1545
DC	05/14/2008	1410	05/14/2008	1426
DC	06/11/2008	1136	06/11/2008	1147
DC	07/09/2008	1021	07/09/2008	1031
DC	08/12/2008	1328	08/12/2008	1338
DC	09/09/2008	1207	09/09/2008	1221
DC	10/07/2008	1140	10/07/2008	1151
DC	11/11/2008	1448	11/11/2008	1500
DC	12/09/2008	1334	12/09/2008	1344
שע	14/09/2008	1334	12/09/2008	1344

7) Associated researchers and projects

As part of the SWMP long-term monitoring program, SAP NERR also monitors Meteorological and Water Quality data which may be correlated with this Nutrient dataset. These data are available from the Research Coordinator or online at http://cdmo.baruch.sc.edu/.

For a complete viewing of associated projects visit the following website and search the collaborators links:

http://gce-lter.marsci.uga.edu/lter/ http://www.uga.edu/marine_advisory/

8) Distribution

NOAA/ERD retains the right to analyze, synthesize and publish summaries of the NERRS System-wide Monitoring Program data. The PI retains the right to be fully credited for having collected and processed the data. Following academic courtesy standards, the PI and NERR site where the data were collected will be contacted and fully acknowledged in any subsequent publications in which any part of the data are used. Manuscripts resulting from this NOAA/OCRM supported research that are produced for publication in open literature, including refereed scientific journals, will acknowledge that the research was conducted under an award from the Estuarine Reserves Division, Office of Ocean and Coastal Resource Management, National Ocean Service, National Oceanic and Atmospheric Administration. 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.

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 http://cdmo.baruch.sc.edu/. Data are available in text tab-delimited format.

II. Physical Structure Descriptors

9) Entry verification

A Lachat QuikChem 8000 FIA+ is used to analyze nutrient concentrations. The instrument is calibrated daily for each parameter to be tested using a series of working standards. Once the calibration run is complete and satisfactory (r >/= 0.99500 up to 1.0000), the samples are set up for analysis. A set of mid-range check standards is used before the sample run, after approximately every 10 samples and at the end of the run to ensure the instrument is in control. The check standards must remain within + or -10% of their original value during the entire run. Also, a blank sample is run and then spiked with each analyte to a known concentration, which must come out within + or -10% as well. Once the run is complete, the raw data is reviewed on the computer attached to the Lachat QuikChem 8000 FIA+ instrument, and the timing is checked to ensure proper integration of sample peaks. Once this is completed, the data is exported via network to another computer. Here the raw file is imported into an Excel spreadsheet and

calculations are performed to obtain the appropriate unit. Orthophosphate values are converted from uM to mg P/L by a conversion factor of 0.03097. Nitrate and nitrite values are converted from uM to mg N/L using a factor of 0.014. Ammonia values are converted from ug N/L to mg N/L by dividing the raw result by 1000. The data file for each month is saved and the results are copied into a comprehensive file with all results. A data quality management (DQM) report is filed with the results. The data was entered and reviewed by Katy Austin Smith, Research Professional II and Lab Manager at the University of Georgia Marine Extension Service.

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 Data Category

Data Category	Parameter	Variable Name	Units of Measure
i) Phosphorus:	*Orthophosphate	PO4F	mg/L as P
ii) Nitrogen:	*Nitrite + Nitrate, Filtered *Nitrite, Filtered *Nitrate, Filtered *Ammonium, Filtered *Dissolved Inorganic Nitrogen	NO23F NO2F NO3F NH4F DIN	mg/L as N mg/L as N mg/L as N mg/L as N mg/L as N
iii) Other Lab F	Parameters: *Chlorophyll a	CHLA_N	μg/L

iv) Field Parameters:

none

Notes:

- 1. Time is coded based on a 2400 hour clock and is referenced to Eastern Standard Time (EST).
- 2. Reserves have the option of measuring either NO2 and NO3 or they may substitute NO23 for individual analyses if they can show that NO2 is a minor component relative to NO3.

11) Measured and Calculated Laboratory Parameters

i) Variables Measured Directly

Nitrogen species: NO2F, NO23F, NH4F

Phosphorus species: PO4F Other: CHLA_N

ii) Computed Variables

NO3F: NO23F-NO2F DIN: NO23F+NH4F

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 UGA Marine Extension Service Laboratory. The MDL is determined as 3 times the standard deviation of a minimum of 7 replicates of a low concentration sample. Table 1 presents the current MDLs; these values are reviewed and revised periodically.

Table 1. Method Detection Limits (MDL) for measured water quality parameters.

Parameter	Variable	Mean Conc.	Std. Dev.	MDL	Dates in use
		mg/L as N or P		mg/L as N or P	
Ammonium	NH4F	0.047	0.001	0.003	Dec.'01 – Dec.'08
Nitrite	NO2F	0.139	0.001	0.004	Jan.'08 – Dec.'08
Nitrite + Nitrate	NO23F	0.126	0.001	0.004	Dec.'01 – Dec.'08
Orthophosphate	PO4F	0.087	0.001	0.002	Dec.'01 – Dec.'08
Chl-a	CHLA			0.0	Dec.'01 – June'07
Chl-a	CHLA	0.6849	0.0053	0.0168	June '07 – Aug. '08
Chl-a	CHLA	0.7987	0.0094	0.0295	Aug. '08 – Dec. '08

13) Laboratory Methods

ii) Parameter: NH4F

OuikChem Method: 31-107-06-1-E

Method Reference: U.S. EPA 1983. USEPA-600/4-79-020. Method 350.1.

Standard Methods 4500-NH₃ H.

Method Descriptor: Samples were filtered with a $0.45~\mu m$ membrane filter and subjected to hypochlorite, which in the presence of phenol, catalytic amounts of nitroprusside and excess hypochlorite, yields indophenol blue, which measured at 630~nm is proportional to the original ammonia concentration.

Preservation Method: Samples filtered and stored frozen (-18 degC).

Holding Time: 2-3 days

iii) Parameter: NO23F

QuikChem Method: 31-107-04-1-C

Method Reference: U.S. EPA 1974. Method 353.2.

Standard Methods 4500-NO₃ F.

Method Descriptor: Samples were filtered with 0.45 um polycarbonate filters. Filtered sample is subjected to cadmium reduction column to reduce nitrate to nitrite. The sample nitrite is then determined by diatizing with sulfanilamide and coupling with N-(1-napthyl)-ethylenediamine dihydrochloride to form a highly colored azo dye which is measured at 520 pm and is proportional to the original nitrate + nitrite concentration. The NO2E

nm and is proportional to the original nitrate + nitrite concentration. The NO2F

concentration (below) is subtracted from this result to give NO3F. Preservation Method: Samples filtered and stored frozen (-18 degC).

Holding Time: 2 weeks

iii) Parameter: NO2F

OuikChem Method: 31-107-04-1-C

Method Reference: U.S. EPA 1974. Method 353.2.

Standard Methods 4500-NO₃ F.

Method Descriptor: Samples were filtered with 0.45 um polycarbonate filters. Nitrite in a filtered sample is measured by closing off the cadmium reduction column so that the nitrate is not converted and the sample follows through the same chemistry as with NO3F to yield the original nitrite concentration.

Preservation Method: Samples filtered and stored frozen (-18 degC).

Holding Time: 1-2 days

iv) Parameter: NO3F

QuikChem Method: 31-107-04-1-C

Method Reference: U.S. EPA 1974. Method 353.2.

Standard Methods 4500-NO₃ F.

Method Descriptor: Nitrate is calculated from NO23F minus NO2F results. Preservation Method: Samples filtered and stored frozen (-18 degC).

Holding Time: 2 weeks

v) Parameter: DIN

Method: DIN is calculated by adding the NH4F and NO23F results together.

vi) Parameter: PO4F

QuikChem Method: 31-115-01-3-A

Method Reference: U.S. EPA 1978. Method 365.1.

Standard Methods 4500-P E.

Method Descriptor: Samples were filtered with 0.45 um polycarbonate filters. Filtered sample is subjected to ammonium molybdate and antimony potassium tartrate under acidic conditions to form a yellow complex. This complex is reduced with ascorbic acid to form a blue complex, which absorbs light at 880 nm. The absorbance is proportional to the concentration of orthophosphate in the sample.

Preservation Method: Samples filtered and stored frozen (-18 degC).

Holding Time: 30 days

vii) Parameter: CHLA

APHA Standard Methods: 10200 H.

Method Reference:

Method Descriptor: Suspended sediment and other material in a water sample is concentrated onto a 47 mm GF/F filter under low vacuum. The sample is stored in a petri dish wrapped in aluminum foil in an airtight plastic bag kept on ice while in the field. The samples are then kept frozen and in the dark until analysis. The acetone extraction method is used to extract the chlorophyll over 2-24 hours and a spectrophotometer is used to obtain readings, which are calculated into a final result.

Preservation Method: Filters are stored frozen (-18 degC).

Holding Time: 28 days

14) **Field and Laboratory QAQC programs** – This section describes field variability, laboratory variability, the use of inter-organizational splits, sample spikes, standards, and cross calibration exercises.

a. Precision

- i. **Field Variability** Field replicates are successive grab samples. These are done in triplicate. Samples are filtered and placed on ice before the next sample is grabbed (usually about 10 minutes between grabs).
- ii. Laboratory Variability Laboratory replicates are done in duplicate.
- iii. Inter-organizational splits –Samples were analyzed by one lab.

b. Accuracy

- i. Sample Spikes A blank sample is spiked with each set for each analyte to obtain a 100 % recovery + or -10 %. One or two sample unknowns are spiked with each set for each analyte to obtain a 100 % recovery + or -20 percent under ideal conditions.
- ii. **Standard Reference Material Analysis** –NERR QA/QC sample analyzed during April 2006; External Standard ("Simple Nutrients" ERA catalog # 739 purchased from Environmental Resource Associates and analyzed with each sample set from August 2008 to December 2008.
- iii. Cross Calibration Exercises None.

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 OAOC 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

Sensor errors

iisoi ciiois	
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

Rec

Record comm	nents
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 WS2 > 10 to 20 knots WS3 > 20 to 30 knots WS4 > 30 to 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.

Diel samples for February 2008 were not collected due to ISCO instrument malfunction.

Samples that have been diluted and rerun are coded CDR. This happens frequently with PO_4 results as those values above the upper limit of the linear range (upper limit 2.2 uM or 0.0682 mg P/L) are diluted, rerun and the appropriate dilution factor applied to the raw data, thus yielding a final result analyzed within the linear range. The following table highlights dilutions that were performed on 2008 samples.

Month	Station ID	Dilution factor
January	Dean Creek (Grab 1 and 2)	10
February	Dean Creek (Grab 1 and 2)	5
March	Dean Creek (Grab 1 and 2)	5
April	Dean Creek (Grab 1 and 2)	2
May	Dean Creek (Grab 1 and 2)	10
June	Cabretta Creek (Grab 1 and 2)	2
June	Dean Creek (Grab 1 and 2)	10
July	Cabretta Creek (Grab 1 and 2)	2
July	Dean Creek (Grab 1 and 2)	10
August	Cabretta Creek (Grab 1 and 2)	2
August	Dean Creek (Grab 1 and 2)	100
August	Hunt Dock (Grab 1 only)	2
September	Cabretta Creek (Grab 1 and 2)	2
September	Dean Creek (Grab 1 and 2)	10
October	Cabretta Creek (Grab 1 and 2)	2

October	Dean Creek (Grab 1 and 2)	5
November	Cabretta Creek (Grab 1 and 2)	2
November	Dean Creek (Grab 1 and 2)	5
December	Cabretta Creek (Grab 1 and 2)	2
December	Dean Creek (Grab 1 and 2)	5