

Chemical Analysis Report

SJRWMD-2020-02-12-01

Florida Department of Environmental Protection
Central Laboratory
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DOH Accreditation E31780

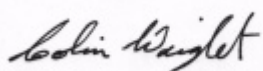
Event Description: **Routine Monitoring for Toxins and Dominant Algal Species (Feb)**
Request ID: **RQ-2020-02-03-77**
Customer: **SJRWMD**
Project ID: **HAB-ROUTNE**

Send Reports to:
St. Johns River Water Management District
St. Johns River Water Management District
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Certified by: Colin Wright, Program Administrator

Date Certified: 14-FEB-2020 08:46



Case Narrative

Unless otherwise noted, all samples included in this report were received in accordance with protocols referenced in Chapter 62-160, Florida Administrative Code (F.A.C.). Results published in this report pertain only to the samples as submitted to, and received by the laboratory. All times in this report are adjusted to the applicable Eastern Time Zone (EST or EDT).

Results for the following analytical group are included in this report: Pesticides.

Scientific notation may be used in reporting very large or small values. Values reported using scientific notation will take the form of the following example: 1.3E+03, which is equivalent to 1.3×10^3 or 1300.

Unless otherwise noted, analytical values for soil and sediment samples are reported on a dry weight basis, and analytical values for waste and tissue samples are reported on a wet weight basis.

Results for TNI accredited tests met requirements established by The NELAC Institute. A double asterisk (**) is used to indicate an analyte/matrix/method for which the laboratory is not TNI accredited by the Florida Department of Health Environmental Laboratory Certification Program or where accreditation for that field of testing is not applicable.

Any significant anomalies or deviations from established protocols are documented in Non-Conformance Reports, which, where appropriate, are included within this analytical report. Additional comments related to specific analytical tests may be included as remarks following the analytical results for each sample. Such comments and remarks are for informational purposes only and are not intended to convey judgement about the usability of the reported data.

A quality control report on the performance of the test method for the submitted samples is included. Uncertainty associated with the analytical results contained in this report can be estimated from the reported quality assurance results and from published quality control acceptance limits for each analytical test. Matrix quality control results (matrix spike recoveries and matrix sample precision) pertain only to the matrix sample tested and do not necessarily reflect test method performance for other samples.

Typical matrix quality control (QC) measurements may include matrix spike recovery, matrix spike duplicate recovery, matrix spike precision and matrix sample precision. Not all matrix QC results may be available or reportable; where they are not an explanation is provided. Typical reasons for unavailable QC results include, but are not limited to, a) insufficient matrix sample to perform some or all QC measurements; b) analyte concentration in the sample replicated was too low for a meaningful measurement of precision and c) analyte concentration in the matrix sample spiked was too high (relative to the amount of analyte spiked) for a meaningful measurement of recovery. Where matrix QC results are unavailable, other method performance metrics (e.g., LCS recovery, LCS precision, surrogate recovery) may be used to assess performance of the method. Comments explaining any missing QC measurements are not intended to convey any adverse conclusions about the quality of the reported data.

Precision is reported as relative percent difference unless otherwise noted.

Quality Control codes as defined below may be used in this report to indicate results that are associated with one or more quality control elements which did not fall within established test method criteria. Such results may be qualified as estimates using a J qualifier as required by 62-160 F.A.C. Explanations are included in the report for any results that were reported as estimates for other reasons.

QC Codes used in this report may include:

- LCS – Recovery for the batch Laboratory Control Sample (LCS) was outside existing control limits;
- MS – Recovery for the batch matrix spike (MS) was outside existing control limits;
- CCV – Recovery for a continuing calibration verification (CCV) standard was outside existing control limits;
- SUR – Recovery of a surrogate (SUR) for associated analytes was outside existing control limits;
- RPD – The precision, measured as relative percent difference (RPD), of batch replicate measurements was outside existing control limits;
- RSD – The precision, measured as relative standard deviation (RSD), of batch replicate measurements was outside existing control limits;
- SMP – Sample - used precision derived from replicate analyses of a sample;

The following data qualifiers are used, where applicable, in this report as specified in 62-160 F.A.C.

- A - Value reported is the mean of two or more determinations.
- B - Results based on colony counts outside the acceptable range.
- I - The reported value is between the laboratory method detection limit and the laboratory practical quantitation limit.
- J - Estimated value and/or the analysis did not meet established quality control criteria.
- K - Actual value is known to be less than value given.
- L - Actual value is known to be greater than value given.
- N - Presumptive evidence of presence of material.
- O - Sampled, but analysis lost or not performed.
- Q - Sample held beyond normal holding time.
- T - Value reported is less than the criterion of detection.
- U - Material was analyzed for but not detected. The reported value is the method detection limit for the sample analyzed.
- V - Analyte was detected in both sample and method blank.
- X - Too few individuals to calculate SCI value.
- Y - The laboratory analysis was from an unpreserved or improperly preserved sample. The data may not be accurate.
- Z - Colonies were too numerous to count (TNTC).

Quality control information from overflow laboratories may not be included in this report. Please refer to the associated report from the overflow laboratory for additional information.

Sample Location: Crescent Lake (Mouth of Dunns Creek)

Collection Date/Time: 02/10/2020 08:49

Field ID: CRESLM

Matrix: W-SURF-FRH

Sample ID	Ref. Method	Component	Result	Code	Units	Batch ID	QC Codes
2156701	EPA 8321B	Anatoxin-a**	0.25	U	ug/L	P378795	
		Cylindrospermopsin**	0.25	U	ug/L	P378795	
		Desmethyl microcystin LR**	0.25	U	ug/L	P378795	
		Microcystin LA**	0.25	U	ug/L	P378795	
		Microcystin LF**	0.25	U	ug/L	P378795	
		Microcystin LR**	0.25	U	ug/L	P378795	
		Microcystin LW**	0.25	U	ug/L	P378795	
		Microcystin LY**	0.25	U	ug/L	P378795	
		Microcystin RR**	0.25	U	ug/L	P378795	
		Microcystin WR**	0.50	U	ug/L	P378795	
		Microcystin YR**	0.25	U	ug/L	P378795	

Quality Assurance Report

Method Blank Results

Reference Method: EPA 8321B
Batch ID: P378795

Component	Result	Code	Units
Anatoxin-a	0.25	U	ug/L
Cylindrospermopsin	0.25	U	ug/L
Desmethyl microcystin LR	0.25	U	ug/L
Microcystin LA	0.25	U	ug/L
Microcystin LF	0.25	U	ug/L
Microcystin LR	0.25	U	ug/L
Microcystin LW	0.25	U	ug/L
Microcystin LY	0.25	U	ug/L
Microcystin RR	0.25	U	ug/L
Microcystin WR	0.50	U	ug/L
Microcystin YR	0.25	U	ug/L

Quality Assurance Report Laboratory Control Sample Accuracy

Reference Method: EPA 8321B
Batch ID: P378795

Component	% Rec.1	% Rec.2	Pass/Fail	Control Limits
Anatoxin-a	95.6		P	40 - 140
Cylindrospermopsin	91.5		P	40 - 140
Desmethyl microcystin LR	82.1		P	40 - 140
Microcystin LA	96.1		P	40 - 140
Microcystin LF	85.4		P	40 - 140
Microcystin LR	82.3		P	40 - 140
Microcystin LW	70.0		P	40 - 140
Microcystin LY	106		P	40 - 140
Microcystin RR	86.2		P	40 - 140
Microcystin WR	87.0		P	40 - 140
Microcystin YR	99.0		P	40 - 140

Quality Assurance Report

Matrix Spike Accuracy

Reference Method: EPA 8321B
Batch ID: P378795

Spiked Sample	Component	% Rec.1	% Rec.2	Pass/Fail	Control Limits
2156080	Anatoxin-a	99.9	101	P/P	40 - 140
2156080	Cylindrospermopsin	84.4	84.1	P/P	40 - 140
2156080	Desmethyl microcystin LR	75.9	86.5	P/P	40 - 140
2156080	Microcystin LA	73.0	82.3	P/P	40 - 140
2156080	Microcystin LF	81.8	69.4	P/P	40 - 140
2156080	Microcystin LR	83.0	91.0	P/P	40 - 140
2156080	Microcystin LW	72.6	77.1	P/P	40 - 140
2156080	Microcystin LY	69.5	79.6	P/P	40 - 140
2156080	Microcystin RR	85.4	90.3	P/P	40 - 140
2156080	Microcystin WR	75.7	81.4	P/P	40 - 140
2156080	Microcystin YR	81.7	80.0	P/P	40 - 140

Quality Assurance Report Precision

Reference Method: EPA 8321B
Batch ID: P378795

Replicated Lab Sample	Component	% RSD/RPD	Sample/Spike/LCS*	Pass/Fail	Control Limits
2156080	Anatoxin-a	0.815	Spike	P	0 - 30
2156080	Cylindrospermopsin	0.425	Spike	P	0 - 30
2156080	Desmethyl microcystin LR	13.0	Spike	P	0 - 30
2156080	Microcystin LA	12.0	Spike	P	0 - 30
2156080	Microcystin LF	16.4	Spike	P	0 - 30
2156080	Microcystin LR	9.19	Spike	P	0 - 30
2156080	Microcystin LW	6.04	Spike	P	0 - 30
2156080	Microcystin LY	13.5	Spike	P	0 - 30
2156080	Microcystin RR	5.55	Spike	P	0 - 30
2156080	Microcystin WR	7.25	Spike	P	0 - 30
2156080	Microcystin YR	2.06	Spike	P	0 - 30

* Sample, spike and/or laboratory control sample precision (LCS) is reported.

Quality Assurance Report Surrogates

Lab Sample ID: 2156701
Field Sample ID: CRESLM

Reference Method	Surrogate	% Rec.	Pass/Fail	Control Limits
EPA 8321B	Carbamazepine-d10	102	P	30 - 165

Quality Assurance Report Calibration Verification

Reference Method: EPA 8321B

Run ID: A97522

Included Lab Sample IDs: 2156701

Component	% Rec.1	% Rec.2	Pass/Fail*	Control Limits
Anatoxin-a	104	90.3	P/P	50 - 160
Cylindrospermopsin	99.1	89.0	P/P	60 - 160
Desmethyl microcystin LR	101	93.7	P/P	60 - 160
Microcystin LA	77.2	93.5	P/P	50 - 160
Microcystin LF	88.9	96.2	P/P	60 - 160
Microcystin LR	84.9	97.4	P/P	60 - 160
Microcystin LW	89.6	97.9	P/P	60 - 160
Microcystin LY	91.6	94.5	P/P	60 - 160
Microcystin RR	96.9	95.0	P/P	60 - 160
Microcystin WR	97.7	93.9	P/P	60 - 160
Microcystin YR	92.0	96.9	P/P	60 - 160

* Pass/Fail determinations are made for each bracketing calibration verification check.

Control limits for initial calibration checks may be different from those for continuing checks, depending on method requirements.
Where they are different, both control limits are provided.

Quality Assurance Report Summary

Ref. Method	Analyte	LCS % Recovery	MS % Recovery		Precision SMP	MS
			LCS	MS		
EPA 8321B	Anatoxin-a	95.6	99.9	101		0.815
	Cylindrospermopsin	91.5	84.4	84.1		0.425
	Desmethyl microcystin LR	82.1	75.9	86.5		13.0
	Microcystin LA	96.1	73.0	82.3		12.0
	Microcystin LF	85.4	81.8	69.4		16.4
	Microcystin LR	82.3	83.0	91.0		9.19
	Microcystin LW	70.0	72.6	77.1		6.04
	Microcystin LY	106	69.5	79.6		13.5
	Microcystin RR	86.2	85.4	90.3		5.55
	Microcystin WR	87.0	75.7	81.4		7.25
	Microcystin YR	99.0	81.7	80.0		2.06

Reference Method Descriptions

Method	Description	Associated Samples
EPA 8321B	Microcystins in water matrices by HPLC/MS/MS	2156701

Preparation and Analysis Log

Ref. Method	Received Date	Prep Date/Time	Prepared By	Analysis Date/Time	Analyzed By	Associated Samples
EPA 8321B	02/12/2020	02/13/2020 09:00	Juyoung Kim	02/13/2020 13:48	Juyoung Kim	2156701