## NEW YORK STATE DEPARTMENT OF ENVIRONMENTAL CONSERVATION

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## Mercury, PCBs, Organochlorine Pesticides, and PFAS Analyses of Fish Collected in 2019 from the Long Island Sound

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## ABSTRACT

The Analytical Services Unit (ASU) at Hale Creek Field Station (HCFS) conducted chemical analyses on a total of 8 fish samples collected from the Long Island Sound. All 8 samples were analyzed for total mercury, total PCBs, selected organochlorine pesticides, and selected per- and polyfluoroalkyl substances (PFAS). Maximum contaminant levels found in the samples were 0.441 µg/g for total mercury, 0.746 µg/g for total PCBs, 0.0741 µg/g for total DDT, 0.0272 µg/g for total chlordanes, 1.00 ng/g for perfluoroundecanoic acid (PFUnA), and 2.44 ng/g for perfluorooctane sulfonamide (PFOSA). Levels were below detection limits for 2,4'-DDE, 2,4'-DDT, heptachlor, heptachlor epoxide, oxychlordane, trans-chlordane, aldrin, photomirex, mirex, hexachlorocyclohexanes, hexachlorobenzene, perfluorohexanoic acid (PFHxA), perfluoroheptanoic acid (PFDA), perfluorodecanoic acid (PFDA), perfluorobutanesulfonic acid (PFBS), perfluorohexane sulfonate (PFHxS) and perfluorooctane sulfonic acid (PFOS).

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#### SAMPLE INFORMATION

This report consists of results of analyses of 8 fish samples collected in 2019 from the Long Island Sound for the Toxic Substance Monitoring Program. The fish collected were 8 Striped Bass (STB). The fish were collected by Kurt Gottschall, David Ellis, and Deb Pacileo from Connecticut Department of Energy and Environmental Protection. Collection records for all samples are attached at the end of this report.

#### LABORATORY METHODS

The ASU analyzed all 8 samples for total mercury, total PCBs, selected organochlorine pesticides, and selected PFAS. The ASU Lab Numbers assigned to the samples were 19-1045-H through 19-1052-H. The ASU program name assigned to the samples was LISound-2019.

<u>Sample preparation.</u> Samples were transported to HCFS where they were stored at -20°C or colder. The samples were prepared for analysis in accordance with HCFS Standard Operating Procedure (SOP) *PrepLab4.* All samples were dissected, ground, and homogenized at HCFS.

<u>Mercury analysis.</u> Samples were analyzed for total mercury in fish tissue by thermal decomposition, amalgamation and atomic absorption spectrophotometry using a Milestone Tri-Cell Direct Mercury Analyzer, DMA-80 [HCFS SOP *HC-405 (Total Mercury)*]. The method is based on EPA method 7473 Mercury in Solids and Solutions by Thermal Decomposition, Amalgamation and Atomic Absorption Spectrophotometry (2007).

**PCB/pesticides analysis.** Samples were analyzed for PCBs and selected organochlorine pesticides by capillary GC-ECD [HCFS SOP *OC1.108 (Organochlorine Residues)*]. At least ten percent of the samples were qualitatively confirmed by capillary GC-MS. Prior to analysis, each sample was freeze-dried and soxhlet-extracted with hexane/acetone (1:1), followed by a florisil cleanup step. All samples were analyzed for three PCB Aroclors (Aroclors 1242 and sum of Aroclors 1254/1260) and 19 organochlorine pesticides and metabolites (4,4'-DDE; 4,4'-DDD; 4,4'-DDT; 2,4'-DDE; 2,4'-DDT; heptachlor; heptachlor epoxide; trans-chlordane; cis-chlordane; trans-nonachlor; cis-nonachlor; oxychlordane; aldrin; photomirex; mirex; HCB; alpha-HCH; beta-HCH; and gamma-HCH). The method is based on *FDA Pesticide Analytical Manual Vol. 1, 3<sup>rd</sup> edition*, Sections 202, 203 and 304.

**PFAS analysis.** Samples were analyzed for selected PFAS by LC/MS/MS using isotopic dilution [HCFS SOP *HC-511 (PFAS)*]. Prior to analysis, each sample was extracted with 0.05 N KOH in methanol followed by ENVI-Carb and SPE cleanup steps. All samples were analyzed quantitatively for 11 PFAS (7 carboxylic acids: PFHxA, PFHpA, PFOA, PFNA, PFDA, PFUnA, and PFDoA; 3 sulfonic acids: PFBS, PFHxS, and PFOS; 1 sulfonamide: PFOSA). Samples were also qualitatively monitored for an additional 11 PFAS (PFTrA, PFTeA, PFPeS, PFHpS, PFNS, PFDS, N-MeFOSAA, N-EtFOSAA, 4:2 FTS, 6:2 FTS, and 8:2 FTS). The method was developed using guidance from the *Department of Defense and Department of Energy consolidated Quality Systems Manual for Environmental Laboratories Version 5.3* and EPA method 533: Determination of Per- and Polyfluoroalkyl Substances in Drinking Water by Isotope Dilution Anion Exchange Solid Phase Extraction and Liquid-Chromatography/Tandem Mass Spectrometry.

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## LABORATORY RESULTS

Results are contained in the following tables:

- Table 1: Sample collection, preparation information, and concentration of Mercury in µg/g wet weight;
- Table 2: Percent moisture, percent lipid, and concentrations of PCBs and DDT in µg/g wet weight;
- Table 3: Concentrations of Chlordane in μg/g wet weight;
- Table 4: Concentrations of PFAS in ng/g wet weight.

In each table, the rows are ordered by lab number.

Concentrations were below the detection limit for 2,4'-DDE, 2,4'-DDT, heptachlor, heptachlor epoxide, oxychlordane, trans-chlordane, aldrin, photomirex, mirex, hexachlorocyclohexanes, hexachlorobenzene, PFHxA, PFHpA, PFOA, PFDA, PFDA, PFBS, PFHxS, and PFOS.

All sample information and results are also contained in file "REP 21-12 (LISound-2019).xlsx", formatted in Excel. General information and a data dictionary for the tables and the Excel file are shown in Appendix A. The quality control procedures and quality control results for these analyses are described in Appendix B. The method detection limit (MDL) for each analyte is listed in Table B1 (Appendix B).

LABNO	TAGNO	SPP	SDATE	LOCATION	PREP	LENMM	WGTG	PROGRAM	Hg
19-1045-H	0997375	STB	20191016	Long Island Sound	SF	931	6770	LISound-2019	0.343
19-1046-H	FA2019044-1	STB	20191008	Long Island Sound	SF	782	3835	LISound-2019	0.224
19-1047-H	FA2019044-2	STB	20191008	Long Island Sound	SF	751	3505	LISound-2019	0.155
19-1048-H	FA2019044-3	STB	20191008	Long Island Sound	SF	856	5860	LISound-2019	0.441
19-1049-H	FA2019044-4	STB	20191008	Long Island Sound	SF	834	4730	LISound-2019	0.401
19-1050-H	FA2019044-5	STB	20191008	Long Island Sound	SF	972	8115	LISound-2019	0.164
19-1051-H	FA2019044-6	STB	20191008	Long Island Sound	SF	900	6490	LISound-2019	0.364
19-1052-H	0997393	STB	20191016	Long Island Sound	SF	1034	12130	LISound-2019	0.353

Note: See Appendix A for general information and a data dictionary for this table.

# Table 2: Percent Moisture, Percent Lipid, and Concentrations of PCBs and DDT in µg/g in Fish Collected the Long Island Sound in 2019

	TACNO	CDD	DOTMOIST			PCB Aroclors	6			DDT and me	etabolites		
LADNU	TAGNO	377	PCTMUIST	PUILPD	AR1242	AR125460	TPCB	PPDDE	PPDDD	PPDDT	OPDDE	OPDDT	TDDT
19-1045-H	0997375	STB	79.58	0.83	0.0158	0.131	0.147	0.0107	-0.002	-0.002	-0.005	-0.005	0.0107
19-1046-H	FA2019044-1	STB	78.23	1.09	0.0156	0.213	0.229	0.0107	-0.002	-0.002	-0.005	-0.005	0.0107
19-1047-H	FA2019044-2	STB	77.93	1.80	0.0199	0.115	0.135	0.00743	-0.002	-0.002	-0.005	-0.005	0.00743
19-1048-H	FA2019044-3	STB	79.82	0.46	0.0336	0.138	0.172	0.0134	-0.002	-0.002	-0.005	-0.005	0.0134
19-1049-H	FA2019044-4	STB	79.18	0.99	0.0832	0.406	0.489	0.0197	0.00439	0.00314	-0.005	-0.005	0.0272
19-1050-H	FA2019044-5	STB	76.35	4.78	0.0854	0.307	0.392	0.0266	0.00853	0.00284	-0.005	-0.005	0.0380
19-1051-H	FA2019044-6	STB	74.75	6.37	0.0798	0.629	0.709	0.0475	0.0104	0.00512	-0.005	-0.005	0.0630
19-1052-H	0997393	STB	70.12	10.49	0.111	0.635	0.746	0.0560	0.0129	0.00518	-0.005	-0.005	0.0741

Note: See Appendix A for general information and a data dictionary for this table.

## Table 3: Concentrations of Chlordane in µg/g in Fish Collected from the Long Island Sound in 2019

	TACNO	CDD			Chlorda	ines		
LADNO	TAGNO	JFF	OXYCHLOR	TRANSCHL	CISCHL	TRANSNON	CISNON	TCHL
19-1045-H	0997375	STB	-0.005	-0.005	-0.005	-0.005	-0.005	0.00000
19-1046-H	FA2019044-1	STB	-0.005	-0.005	-0.005	-0.005	-0.005	0.00000
19-1047-H	FA2019044-2	STB	-0.005	-0.005	-0.005	-0.005	-0.005	0.00000
19-1048-H	FA2019044-3	STB	-0.005	-0.005	-0.005	-0.005	-0.005	0.00000
19-1049-H	FA2019044-4	STB	-0.005	-0.005	-0.005	-0.005	-0.005	0.00000
19-1050-H	FA2019044-5	STB	-0.005	-0.005	-0.005	-0.005	-0.005	0.00000
19-1051-H	FA2019044-6	STB	-0.005	-0.005	0.00697	0.00866	0.00546	0.0211
19-1052-H	0997393	STB	-0.005	-0.005	0.00912	0.0116	0.00649	0.0272

Note: See Appendix A for general information and a data dictionary for this table.

## Table 4: Concentration of PFAS in ng/g in Fish Collected from the Long Island Sound in 2019

	TACNO	CDD						PFAS					
LADINU	TAGNO	377	PFHxA	PFHpA	PFOA	PFNA	PFDA	PFUnA	PFDoA	PFBS	PFHxS	PFOS	PFOSA
19-1045-H	0997375	STB	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-2.00	-2.00	-2.00	-2.00
19-1046-H	FA2019044-1	STB	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-2.00	-2.00	-2.00	-2.00
19-1047-H	FA2019044-2	STB	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-2.00	-2.00	-2.00	-2.00
19-1048-H	FA2019044-3	STB	-1.00	-1.00	-1.00	-1.00	-1.00	1.00	-1.00	-2.00	-2.00	-2.00	-2.00
19-1049-H	FA2019044-4	STB	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-2.00	-2.00	-2.00	2.44
19-1050-H	FA2019044-5	STB	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-2.00	-2.00	-2.00	-2.00
19-1051-H	FA2019044-6	STB	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-2.00	-2.00	-2.00	-2.00
19-1052-H	0997393	STB	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-1.00	-2.00	-2.00	-2.00	-2.00

Note: See Appendix A for general information and a data dictionary for this table.

## APPENDIX A

General information for using tables and electronic file: "REP 21-12 (LISound-2019).xlsx"

- 1. Chemical concentrations are reported in µg/g (ppm) and ng/g (ppb) wet weight.
- 2. The results are reported to no more than three significant figures.
- 3. A negative concentration indicates the concentration was below the MDL. The number following the negative sign is the MDL.

## Data dictionary for tables and electronic file: "REP 21-12 (LISound-2019).xlsx"

- 1. LABNO unique sample lab number assigned at Hale Creek Field Station (character)
- 2. TAGNO sample identifier assigned at time of collection and contained in collection records (character)
- 3. SPP species code; STB=Striped Bass (character)
- 4. SDATE date sample was collected; format is YYYYMMDD (numeric)
- 5. LOCATION location where sample was collected (character)
- 6. AGE age of fish in years, if determined (numeric)
- 7. SEX sex of fish, if determined; M=male; F=female (character)
- 8. PREP preparation method; SF=standard fillet, W=whole fish; W-HV=whole fish minus the head and viscera (character)
- 9. LENMM fish length in mm; mean length in mm if sample is composite (numeric)
- 10. WGTG fish weight in g; total weight in g if sample is composite (numeric)
- 11. PROGRAM program name assigned by Hale Creek Field Station (character)
- 12. MAXLEN maximum fish length in mm if sample is composite (numeric)
- 13. MINLEN minimum fish length in mm if sample is composite (numeric)
- 14. SDLEN standard deviation of fish length in mm if sample is composite (numeric)
- 15. MAXWGT maximum fish weight in g if sample is composite (numeric)
- 16. MINWGT minimum fish weight in g if sample is composite (numeric)
- 17. SDWGT standard deviation of fish weight in g if sample is composite (numeric)
- 18. NOANLY number of individuals in sample; if NOANLY is greater than 1, then sample is composite (numeric)
- 19. PCTMOIST percent moisture in sample (numeric)
- 20. PCTLPD percent lipid in sample (numeric)
- 21. Hg mercury (numeric)
- 22. AR1242 Aroclor 1242 (numeric)
- 23. AR125460 sum of Aroclors 1254 and 1260 (numeric)
- 24. TPCB total PCBs; sum of AR1242 and AR125460 (numeric)
- 25. PPDDE 4,4'-DDE (numeric)
- 26. PPDDD 4,4'-DDD (numeric)
- 27. PPDDT 4,4'-DDT (numeric)
- 28. OPDDE 2,4'-DDE (numeric)
- 29. OPDDT 2,4'-DDT (numeric)
- 30. TDDT total DDT; sum of PPDDE, PPDDD, PPDDT, OPDDE and OPDDT (numeric)
- 31. HEPT heptachlor (numeric)
- 32. HEPTEPOX heptachlor epoxide (numeric)
- 33. THEPT total heptachlor; sum of HEPT and HEPTEPOX (numeric)
- 34. OXYCHLOR oxychlordane (numeric)
- 35. TRANSCHL trans-chlordane (numeric)
- 36. CISCHL *cis*-chlordane (numeric)
- 37. TRANSNON *trans*-nonachlor (numeric)
- 38. CISNON *cis*-nonachlor (numeric)
- 39. TCHL total chlordanes; sum of OXYCHLOR, TRANSCHL, CISCHL, TRANSNON and CISNON (numeric)
- 40. ALDRIN aldrin (numeric)
- 41. MIREX mirex (numeric)
- 42. PHOMIREX photomirex (numeric)
- 43. TMIREX total mirex; sum of MIREX and PHOMIREX (numeric)
- 44. AHCH α-hexachlorocyclohexane; α-BHC; α-benzene hexachloride (numeric)
- 45. BHCH β-hexachlorocyclohexane; β-BHC; β-benzene hexachloride (numeric)
- 46. GHCH γ-hexachlorocyclohexane; γ-BHC; γ-benzene hexachloride; lindane (numeric)
- 47. THCH total HCH; sum of AHCH, BHCH and GHCH (numeric)
- 48. HCB hexachlorobenzene (numeric)
- 49. PFHxA Perfluorohexanoic acid (numeric)
- 50. PFHpA Perfluoroheptanoic acid (numeric)
- 51. PFOA Perfluorooctanoic acid (numeric)
- 52. PFNA Perfluorononanoic acid (numeric)
- 53. PFDA Perfluorodecanoic acid (numeric)
- 54. PFUnA Perfluoroundecanoic acid (numeric)

- 55.
- 56.
- PFDoA Perfluorododecanoic acid (numeric) PFBS Perfluorobutanesulfonic acid (numeric) PFHxS Perfluorohexanesulfonic acid (numeric) PFOS Perfluorooctanesulfonic acid (numeric) PFOSA Perfluorooctane sulfonamide (numeric) 57.
- 58.
- 59.

## APPENDIX B

#### Quality control for mercury

To determine mercury concentration, each sample was analyzed twice (two separate aliquots of the same homogenate) and the average was reported.

The quality control for mercury included analyses of, at minimum, one reference material sample, one laboratory duplicate, and one method blank for every 20 samples. For the reported analyses, there was one method blank, four reference material samples, and one duplicate sample. The reference materials were one SRM 2976, two SRM 1947, and two DORM-4 Dogfish Muscle from NRC, Canada. For each laboratory duplicate analysis, four aliquots of the same homogenate were analyzed; the first and second aliquots were averaged and the third and fourth aliquots were averaged, and the absolute value of the relative percent difference ([RPD]) of the two averages was determined. The reference material sample and laboratory duplicate results were used to determine accuracy and precision, respectively, of the fish tissue sample results. The procedure blank (laboratory water used during the analysis procedure) was analyzed to determine potential contamination of fish tissue samples.

Criteria for control limits for mercury were based on recommended control limits in *Guidance for Assessing Chemical Contaminant Data for Use in Fish Advisories, Volume 1, 3<sup>rd</sup> edition* (USEPA Office of Water, November 2000) with more stringent modifications as recommended by the instrument manufacturer. Control limits for accuracy were percent recovery = 90-110 percent. The control limit for precision was the relative percent difference (RPD) of laboratory duplicate analyses  $\leq$  20 percent. The MDL was used to assess potential contamination. The statistically derived MDL was 0.004 µg/g Hg wet weight.

Total mercury in the method blank was below the MDL. The percent recovery of total mercury from the reference material was 100 percent for DORM-4; 100 percent for SRM 2976, and 98.8 percent for SRM 1947. The RPD for the laboratory duplicate was 10.5 percent.

#### Quality control for PCBs/organochlorine pesticides

To better assess the overall accuracy and precision of the large number of organic analytes that are measured, a quality control summary is presented for the analysis dates of August 06, 2020 through October 14, 2020 includes the analyses of fish from the Long Island Sound in 2019. The quality control for this period included analyses of seven matrix spikes, six reference materials (six HRM), seven laboratory duplicates, and six method blanks. One matrix spike, reference material, laboratory duplicate, and method blank were analyzed for every 20 samples. The matrix spikes, reference material samples, and laboratory duplicates were used to determine accuracy and precision of the fish tissue sample results. The method blanks (solvent carried through the entire extraction, clean-up and analysis procedure) were used to determine potential contamination of the fish tissue samples.

Criteria for control limits were based on recommended control limits in Guidance for Assessing Chemical Contaminant Data for Use in Fish Advisories, Volume 1, 3rd edition (USEPA Office of Water, November 2000). Control limits for accuracy were percent recovery = 50-150 percent. The control limit for precision was relative standard deviation (RSD)  $\leq 50$  percent. The MDL was used to assess potential contamination.

The control limit for accuracy was determined to be exceeded for an analyte in the study if the mean percent recovery from the matrix spikes or reference material was outside 50-150 percent (see Table B1).

The control limit for precision was determined to be exceeded for an analyte in the study if the RSD of any of the following measures was greater than 50 percent (see Table B1):

- RSD of replicate analyses of matrix spikes or
- RSD of replicate analyses of the reference material or
- mean RSD of laboratory duplicate.

All analytes in the method blanks were below the MDL. The MDLs for the analytes are listed in Table B1.

## **Quality control for PFAS**

The quality control for PFAS included analyses of, at minimum, one reference material sample, one laboratory control sample, one laboratory duplicate, and one method blank for every extraction batch of up to 20 samples. For the reported analyses, there were one method blank, one reference material sample, one laboratory control sample and one duplicate sample. The reference material was one SRM 1947. The reference material samples, laboratory control samples, and laboratory duplicate results were used to determine accuracy and precision of the fish tissue sample results. The method blanks (laboratory water used during the analysis procedure) were analyzed to determine potential contamination of fish tissue samples. Criteria for control limits for PFAS were based on recommended control limits in EPA method 533: Determination of Per- and Polyfluoroalkyl Substances in Drinking Water by Isotope Dilution Anion Exchange Solid Phase Extraction and Liquid-Chromatography/Tandem Mass

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Spectrometry. Control limits for accuracy were percent recovery = 70-130 percent. The control limit for precision was the relative percent difference (RPD) of laboratory duplicate analyses  $\leq 30$  percent. The MDL was used to assess potential contamination.

The control limit for accuracy was determined to be exceeded for an analyte in the study if the percent recovery from the laboratory control sample or reference material was outside 70-130 percent (see Table B2).

The control limit for precision was determined to be exceeded for an analyte in the study if the RSD of any of the following measures was greater than 30 percent (see Table B2).

All analytes in the method blanks were below the MDL. The MDLs for the analytes are listed in Table B2.

### Summary of quality control

All quality assurance was within control limits for accuracy, precision, and potential contamination in ASU Report 21-12.

ANALYTE	MATRIX	SPIKE	REFEF MATE	RENCE RIAL *	LABOR DUPLIC	ATORY ATES **	MDL
	MEAN %R	RSD (%)	MEAN %R	RSD (%)	# of PAIRS	MEAN RSD %	(ng/g)
Aroclor 1242	106	4.72	-	-	6	6.05	10
Aroclor 1254/1260	108	2.78	-	-	6	7.51	30
SUM Aroclor			114	5.02			-
4,4'-DDE	111	3.98	-	-	7	7.18	2
4,4'-DDD	111	5.42	-	-	6	9.44	2
4,4'-DDT	109	6.29	-	-	6	11.5	2
2,4'-DDE	116	3.84	-	-	-	-	5
2,4'-DDT	111	5.58	-	-	-	-	5
Heptachlor	104	9.38	-	-	-	-	5
Heptachlor epoxide	104	3.36	-	-	-	-	5
trans-Chlordane	103	5.64	-	-	-	-	5
cis-Chlordane	107	4.20	-	-	-	-	5
trans-Nonachlor	107	5.53	-	-	1	7.63	5
cis-Nonachlor	96.6	4.25	-	-	-	-	5
Oxychlorodane	105	3.69	-	-	-	-	5
Aldrin	84.5	9.22	-	-	-	-	5
Photomirex	105	4.04	-	-	-	-	5
Mirex	101	4.36	-	-	-	-	2
alpha-HCH	87.1	19.1	-	-	-	-	5
beta-HCH	97.5	16.10	-	-	-	-	5
gamma-HCH	91.3	10.37	-	-	-	-	5
НСВ	79.2	11.2	-	-	1	9.20	2

\*Reference material for SUM Aroclor was HRM (N=6). \*\*Laboratory duplicate RSDs were only used to calculate a mean RSD when the result for each sample in the pair was greater than the MDL.

 Table B2: Percent Recovery, Precision, and MDLs of Per- and Polyfluoroalkyl Substances in One Laboratory

 Control Spike, One Reference Material Sample, and One Pairs of Laboratory Duplicate Analyzed at

 Hale Creek Field Station for Fish Collected from the Long Island Sound in 2019.

ANALYTE	LABOR/ CONT SAM	ATORY ROL PLE	REFEF MATE	RENCE RIAL *	LABOR DUPLIC	ATORY ATES **	MDL
	MEAN %R	RSD (%)	MEAN %R	RSD (%)	# of PAIRS	MEAN RSD %	(ng/g)
PFHxA	122%	-	-	-	-	-	1
PFHpA	121%	-	-	-	-	-	1
PFOA	119%	-	-	-	-	-	1
PFNA	123%	-	-	-	-	-	1
PFDA	126%	-	-	-	-	-	1
PFUnA	125%	-	-	-	-	-	1
PFDoA	124%	-	-	-	-	-	1
PFBS	111%	-	-	-	-	-	2
PFHxS	80.1%	-	-	-	-	-	2
PFOS	102%	-	89.2%	-	-	-	2
PFOSA	102%	-	-	-	-	-	2

\*Reference material for PFOS was SRM 1947 (N=1).

\*\*Laboratory duplicate RSDs were only used to calculate a mean RSD when the result for each sample in the pair was greater than the MDL.

# APPENDIX C: Chain of Custody and Collection Records

NEW YORK STATE D	EPARTMENT OF ENVIRONME CHAIN OF CUSTODY	NTAL CONSERVATION
I, Kunt Cester In 11 (Print Name)	, of <u>CT DEEP 333 Fer</u> (Print Busi	ness Address)
following on 10/10/19	, 20 from Island	Sam
in the vicinity of ALZ of	Math have the box 5.	(Water Body) k. c. 3 30
Town of	(Landmark. Village, Road, etc.) L	16 7230.060 Suffel
Item(s) 1 Staped	B.55 - tag # 0997375	
Said sample(s) were in my posse	ession and handled according to stand	1
collection. The sample(s) were p Environmental Conservation of	placed in the custody of a representative	of the New York State Department
ermt	At Mente: 21	_, 20_19
Signatur	e	<u>2/14/2020</u> Date
1, Coutlin Crais	, received the above mentione	ed sample(s) on the data and if it
and assigned identification number	er(s) 0997375	to the sample(s) of the sample(s).
have recorded pertinent data for t	he sample(s) on the attached collection	records. The sample(s) remains t i
my custody until subsequently tra	insferred, prepared or shipped at times a	and on dates as attack by build
11 1 0	0	/ / .
Signat	ture)	Date
SECOND RECIPIENT (Print Name)	TIME & DATE	2 uto
Megan Barrow	2/5/2020 8 -4	PURPOSE OF TRANSFER
SIGNATURE / M	UNIT 3.57	PCB testing
THIRD RECEIVENT	Diadremous	
(Print Name)	TIME & DATE	PURPOSE OF TRANSFER
SIGNATURE	LINDT	
	UNIT	
FOURTH RECIPIENT (Print Name)	TIME & DATE	N ID DO NOT
		PURPOSE OF TRANSFER
SIGNATURE	UNIT	-
RECEIVED INT ADDR		
RECEIVED IN LABORATORY BY (Print Name)	TIME & DATE	REMARKS
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	11 - 50	
LOGGED IN BY (Print Name)	HCFS	
LOOGED IN BY (Print Name)	HCFS TIME & DATE	ACCESSION NUMBERS
LOOGED IN BY (Print Name)	HCFS TIME & DATE 1.25 pm 2-10-2000	ACCESSION NUMBERS
LOGGED IN BY (Print Name)	HCFS TIME & DATE 1.25 pm 2-10-2070 UNIT ILCCC	ACCESSION NUMBERS 19.1045-H

LACE VY	Method: DElectrofis	hing □Gill net ng □Other	ting Tra	p netting Arawling Notes (SY	g 🗆 Seinir WFDB sur	ng 🗆 Angli vey number	ng ⊡Other ): <i>FA201</i>	9051	s.4c 0330	
R LAB USE	COLLECTION OR TAG NO.	SPECIES	DATE		AGE	SEX &/OR REPROD. CONDIT	LENGTH (mm)	(S)	REMARKS	113-31
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n an							ge of protection	-		352
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ж. <sup>10</sup>							 0 5-524 5 21 - 3			
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ASU Report 21-12

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NEW YORK STATE DEPAI	RTMENT OF ENVIRONMENTA CHAIN OF CUSTODY	AL CONSERVATION
I, Da viet) 5/11:5, (Print Name)	of CT DSS C 333 Fare R (Print Business	Address)
following on <u>Cat</u> Sth, (Date)	2019 from Levy Tisterel	Service (
in the vicinity of Marth, of	(Landmark, Village, Road, etc.)	1
Town of Cla Lyme	in Nrul Low	CT County.
Item(s)	Stripped Bass - tags	FA20190441 -> 6
Said sample(s) were in my possession collection. The sample(s) were place	n and handled according to standard p d in the custody of a representative of	rocedures provided to me prior to the New York State Department of
Environmental Conservation on	November 21,	20_14
Signature	lln	2/14/2020 Date
1. Coutlin Craw	, received the above mentioned a	sample(s) on the date specified
and assigned identification number(s	FA2019044-1->6	to the sample(s). I
have recorded pertinent data for the s	ample(s) on the attached collection rec	cords. The sample(s) remained in
my custody until subsequently transf	erred, prepared or shipped at times and	l on dates as attested to below.
Signature	5	21/19 Date
SECOND RECIPIEN'I' (Print Name)	TIME & DATE	PURPOSE OF TRANSFER
SIGNATURE .	2/5/7020 3:54 UNIT	Pers testing
THIRD BECIPIENT (Print Name)	Diadiome S	PURPOSE OF TRANSFER
SIGNATURE	UNIT	
FOURTH RECIPIENT (Print Name)	TIME & DATE	PURPOSE OF TRANSFER
SIGNATURE	UNIT	14 
RECEIVED IN LABORATORY BY (Print Name)	TIME & DATE	REMARKS
SIGNATURE C	UNIT UNIT	
LOGGED IN BY (Print Name)	TIME & DATE	ACCESSION NUMBERS
Chloc Armato	1:27pm 2-10-2020	19-1046-н>
SIGNATURE	HCFS.	19-1051-4
richter: revised 21 April 2014; becker: 23 March	2017	

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Region		4: 15.34 1240 pri	REMARKS	Toker lengty - 78-2	TL = V 751	TL - 856	71- 534	225 - 72	71 - 900										
DEC	jr	19044 SI	WEIGHT	3835	3505	5860	4730	8115	6490 3										
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D D D D	ing DAng	urvey numbe	SEX &/OR REPROD. CONDIT			- Schwarten 1977	2			· · · · ·	No.		11 - 					3/97 	n finis n
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NEW YORK	hing DGill n	lg Other	SPECIES	578	STR	STR	578	378	s T.S		n n National Line					1 4415	1	- 	17; becker: 3/23/17
d Site Name Zande	Method: DElectrofis	on Method: Freezir	COLLECTION OR TAG NO.	1 - hhobicery	FAZOPO44 - 2	FARMPOHH - 3	FARDISO44-4	- HHOLIOEUL	Hargouy - 6		N. N.								2011, 5/7/15, 10/4/16, 3/20/
Project and Collections	Sampling	Preservatic	FOR LAB USE ONLY- LAB ENTRY NO.	19-1046-H	H-LHOI-PI	H-8401-61	H-PHOI-P1	H-0501-6	H-1501-61									1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1	richter: revised

NEW YORK STATE DEPAR	TMENT OF ENVIRONMENTA) CHAIN OF CUSTODY	L CONSERVATION
I, Kunt Castistall, o	f CT WS P 333 Frank (Print Business)Ac	101.1Lanc_LT collected the
following on <u>Oct</u> 16, 2 (Date)	019 from Lorry Island	Body)
in the vicinity of wast of	(Landmark, Village. Road, etc.)	Lat/Lon
Town of	, in <u>Suff</u> o	117 7734,970 County.
Item(s) ) JAnpal Bes	55 - tag # 0997393	
Said sample(s) were in my possession collection. The sample(s) were placed	and handled according to standard pro- in the custody of a representative of t	beedures provided to me prior to the New York State Department of
Environmental Conservation on	Nogen bis 21, 2	0 <u>19</u>
Signature		Date
1. Caitlin Cary	, received the above mentioned s	ample(s) on the date specified
and assigned identification number(s)	0997393	to the sample(s). I
have recorded pertinent data for the sa	imple(s) on the attached collection rec	ords. The sample(s) remained in
my custody until subsequently transfe	rred, prepared or snipped at times and	
Signature		Date
SECOND RECIPIENT (Print Name)	TIME & DATE	PURPOSE OF TRANSFER
Megan Barrow	2/5/2020 8:54	PCB testiny
SIGNATURE " C Burn	Diadromors	
THIRD RECIPIENT (Print Name)	TIME & DATE	PURPOSE OF TRANSFER
SIGNATURE	UNIT	
FOURTH RECIPIENT (Print Name)	TIME & DATE	PURPOSE OF TRANSFER
SIGNATURE	UNIT	
RECEIVED IN LABORATORY BY (Print Name)	TIME & DATE	REMARKS
SIGNATURE	UNIT HCFS	
LOGGED IN BY (Print Name)	TIME & DATE	ACCESSION NUMBERS
SIGNATURE SIGNATURE	UNIT 415-2020	1-105 2 11
richter, revised 21 April 2014; becker: 23 March	HCFS	

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IFE	ng ⊔Angl rvey numbe	SEX &/OR REPROD. CONDIT				5 5 						2 2 2 2					e second en	7-31-9 
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SION OF FISH AND SION OF FISH AND SH COLLECTION R	netting Arawling Notes (SW	TOCATION LOCATION	234.973		-955 1975	2) ·	is selected a fra Gal			2.34) - 143	0 (1182 9 (39) 3 			i di li noi in A			etus etus	
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NEW YORK:	ning 🛛 Gill net 🖉	SPECIES	STR															/17: becker: 3/23/17
Site Name	Aethod: DElectrofish n Method: Efreezin	COLLECTION OR TAG NO.	567393				2											2011 5/7/15 10/4/16 3/20
Project and	Sampling A Preservatio	FOR LAB USE ONLY- LAB ENTPV NO	19-1052-H			· ·	~		. Al				in the second		1-10			richter revieed