

Session 2: Analytical methods for PFAS in environmental media

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PFAS Science Webinars for EPA Region 1 and State & Tribal Partners

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Outline

- Types of analytical and standard methods
- Drinking Water Sample Methods
 - Method 537
 - Method 533
- Non-Drinking Water Sample Methods
 - SW-846 Method 8327 Direct Injection
 - CWA/SW846 Method—Isotope Dilution
- PFAS Analysis in Marine Waters
- PFAS Analysis in Fish Tissue
- Total Organofluorine Analysis using Combustion Ion Chromatography (TOF)
- Total Organic Precursors (TOP)
- Summary of EPA PFAS Methods as of July 2020



Methods – Sampling

Guidance to avoid cross contamination in sampling

- No teflon
- Avoid contact with clothes, materials containing PFAS (e.g. some food wrappers)
- See <u>PFAS Analytical Methods Development and Sampling website</u>
 - Sampling guidance from states (e.g. MI)
 - Interstate Technology and Regulatory Council Fact Sheet: Site Characterization Considerations, Sampling Precautions, and Laboratory Analytical Methods for PFAS
- PFAS Quality Assurance Plan and Data Review technical brief



Non-Targeted Analysis

Targeted methods are methods which are applicable to a specific defined set of known analytes

- Analytical standards exist for quantitation
- Method only 'sees' analytes on the targeted list will not measure others
- 'One and done' once the analysis is complete, can't look for other analytes

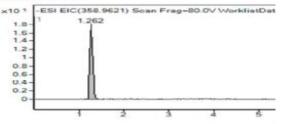
Non-targeted methods involve the use of High Resolution Mass Spectrometry (HRMS) capable of identifying all analytes in a sample, known and unknown

- Can quantitate those for which laboratory standards exist, otherwise may semi-quantitate based on known, structurally similar analytes
- Can screen for lists of known suspects, can discover new/unknown analytes
- Can store the HRMS data and go back later to look for analytes which were unidentified at the time of analysis, but which later become known

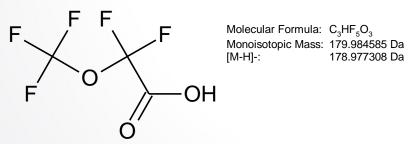
Targeted vs Non-Targeted

Explore unknown compounds using high resolution mass spectrometry. Identify a peak in a chromatogram and to ultimately predict the identity of this unknown.

Mass spectrometer assigns a high resolution mass for peaks observed in the chromatogram



- Software calculates the exact number and type of atoms needed to achieve the measured mass
- ↓ Fragmentation experiments allow determination of most likely structure:



- Using mass, formula and structure, identity can be assigned by searching against databases of known compounds
- ↓ Compare peak to commercial to confirm identification if possible

©EPA Types of Analytical Methods

Three broad classes of methods:

- EPA Standard Methods
 - Methods which have been through a multi-lab validation following a particular rulemaking or guidance effort and are available to support Agency regulatory or guidance activities

Research Methods

- Methods which have been developed by an EPA ORD laboratory for research purposes; QA'ed and peer reviewed via publication, but not multi-lab validated, not considered EPA Standard Methods
- Developmental Methods
 - Methods which are currently undergoing research, development and testing; might become Standard Methods or Research Methods

SEPA Types of Standard Methods

Three broad categories of EPA Standard Methods related to water:

- Safe Drinking Water Act Methods
- <u>Clean Water Act Methods</u>
- SW846 Methods

These are generally targeted methods for solids and water.

Drinking Water Method 537: Revision I

- Update: External lab validation for additional analytes by 537
 - Perfluoro-2-propoxypropanoic acid (GenX chemical HFPO-DA, CAS 13252-13-6)
 - Potassium 9-chlorohexadecafluoro-3-oxanone-1-sulfonate (9CI-PF3ONS, CAS 73606-19-6)
 - Potassium 11-chloroeicosafluoro-3-oxaundecane-1-sulfonate (11Cl-PF3OUdS, CAS 83329-89-9)
 - Sodium dodecafluoro-3H-4,8-dioxanonate (ADONA, CAS 958445-44-8)
- Incorporated clarifications issued in <u>EPA Technical Advisory-</u> <u>Laboratory Analysis of Drinking Water Samples for</u> <u>Perfluorooctanoic Acid (PFOA) Using EPA Method 537 Rev. 1.1</u>
- <u>Final published method</u> (November 2018)

SEPA

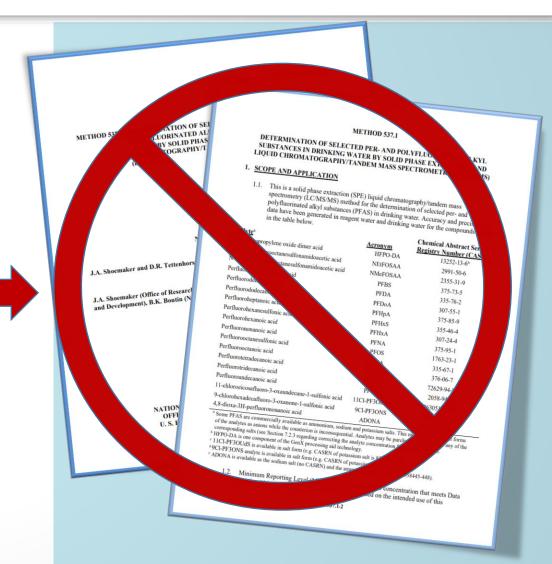
 LC/MS/MS with internal standards. Single lab lowest concentration minimum reporting levels (LCMRLs) range from 0.53-6.3 ng/L

	CPA Document #: EPA/600/R-18/352		
METHOD 537.1 DETERMINATION OF SE POLYFLUORINATED AL WATER BY SOLID PHAS CHROMATOGRAPHY/T (LC/MS/MS)	METHOD 537.1 DETERMINATION OF SELECTED PER- AND POLYFLUORINATED ALKYL SUBSTANCES IN DRINKING WATER BY SOLID PHASE EXTRACTION AND LIQUID CHROMATOGRAPHIV/TANDEM MASS SPECTROMETRY (LCMSMS) SCOPE AND APPLICATION 1. This is a solid phase extraction (SPE) liquid chromatography/tandem mass spectrometry (LCMSMS) method for the determination of selected per- and polyfluorinated alkyl substances (PFAS) in drinking water. Accuracy and precision data have been generated in reagent water and drinking water for the compounds listed in the table below.		
		and drinking	water for the compounds listed
. 1	N-ethyl perfluorooctanesulfonamidoacetic acid N-methyl perfluorooctanesulfonamidoacetic acid	Acronym HFPO-DA	Chemical Abstract Services Registry Number (CASRN) 13252-13-6 ^b
J.A. Shoemaker and D.R. Tettenhors	Perfluorobutanonale	NEtFOSAA	2991-50-6
J.A. Sheen	Perfluorodecanoic and	NMeFOSAA	2355-31-9
c Decourel	Perfluorododecanoic acid	PFBS	375-73-5
J.A. Shoemaker (Office of Researc) and Development), B.K. Boutin (N	Perfluoroheptanoic acid	PFDA	335-76-2
	Perfluorohexanesulfonic acid	PFDoA	307-55-1
	Perfluorohexanoic acid	PFHpA	375-85-9
	Perfluorononanoic acid	PFHxS	
	Perfluorooctanesulfonic acid	PFHXA	355-46-4
	Perfluorooctanoic acid	PFNA	307-24-4
	Perfluorotetradecanoic acid	PFOS	375-95-1
	Perfluorotridecanoic acid	PFOA	1763-23-1
	Perfluoroundecanoic acid	PFTA	335-67-1
	11-chlorogicage 0	PFTrDA	376-06-7
	11-chloroeicosafluoro-3-oxaundecane-1-sulfonic acid 9-chlorohexadecafluoro-3-oxaundecane-1-sulfonic acid	PFUnA	72629-94-8
	4.8-diova 211 - Continone-1-stil foni-	11CI-PF3OUdS	2058-94-8
NATION OFFI		9CI-PF3ONS	763051-92-9*
U.S.I	of d	ADONA	/36426-58-1d
	¹ Some PFA3 are commercially svailable as ammenium, sodim of the analytes as anione shall be counterion in income in other analytes as anione shall be counterion in income in our responding as invalidable the Gran Yunescang of the PIEPO-DA is one composited in sait form (e.g. CASRN of pot 9CC-PF30NS analyte in available in a sait form (e.g. CASRN of ADONA is available as the sodium sait (no CASRN) and the an ADONA is available as the sodium sait (no CASRN) and the Quality Objectives (DQOs) that are develop S37.12	nology, tssium salt is 83329-89-9). potassium salt is 73606-19- nmonium salt (CASPA).	e salt content).

Drinking Water Method 537: "Modified"

- Method 537 is often modified by analytical laboratories for use on non-drinking water samples
- If modifications are made that are not explicitly listed in 537 or 537 Revision 1, the method is not considered 537 by EPA
- The most common modification is inclusion of isotope dilution

EPA





Drinking Water Method 533

Solid phase extraction/isotope dilution method targeting PFAS <C12

- Method 537 generally performs poorly for C4 compounds (e.g. PFBA, PFBS)
- Solid phase extraction, LC/MS/MS, Isotope dilution
- Will allow EPA to consider additional PFAS for monitoring under the fifth Unregulated Contaminant Monitoring Rule
- Released December 2019
 - <u>Analytical Methods Developed by EPA for Analysis of Unregulated</u> <u>Contaminants</u>



Drinking Water Method 533

Method 533	Both Methods	Method 537.1	
1H, 1H, 2H, 2H-perfluorodecane sulfonic acid (8:2 FTS)	11-chloroeicosafluoro-3-oxaundecane-1-sulfonic acid (11Cl-PF3OUdS)	N-ethyl perfluorooctanesulfonamidoacetic acid (NEtFOSAA)	
1H, 1H, 2H, 2H- perfluorohexane sulfonic acid (4:2 FTS)	9-chlorohexadecafluoro-3-oxanone-1-sulfonic acid (9Cl-PF3ONS)	N-methyl perfluorooctanesulfonamidoacetic acid (NMeFOSAA)	
1H, 1H, 2H, 2H-perfluorooctane sulfonic acid (6:2 FTS)	4,8-dioxa-3H-perfluorononanoic acid (ADONA) ³	Perfluorotetradecanoic acid (PFTA)	
Nonafluoro-3,6-dioxaheptanoic acid (NFDHA)	Hexafluoropropylene oxide dimer acid (HFPO-DA)	Perfluorotridecanoic acid (PFTrDA)	
Perfluoro (2-ethoxyethane) sulfonic acid (PFEESA)	Perfluorodecanoic acid (PFDA)		
Perfluoro-3-methoxypropanoic acid (PFMPA)	Perfluorododecanoic acid (PFDoA)		
Perfluoro-4-methoxybutanoic acid (PFMBA)	Perfluorohexanoic acid (PFHxA)		
Perfluorobutanoic acid (PFBA)	Perfluoroundecanoic acid (PFUnA)		
Perfluoroheptanesulfonic acid (PFHpS)	Perfluorobutanesulfonic acid (PFBS)		
Perfluoropentanesulfonic acid (PFPeS)	Perfluoroheptanoic acid (PFHpA)		
Perfluoropentanoic acid (PFPeA)	Perfluorohexanesulfonic acid (PFHxS)		
	Perfluorononanoic acid (PFNA)		
	Perfluorooctanoic acid (PFOA)		
	Perfluorooctanesulfonic acid (PFOS)		



Non-Drinking Water Sample Methods: SW-846 Method 8327—Direct Injection

Non-drinking water aqueous matrices:

- Groundwater
- Surface water
- Wastewater

Find a balance among sensitivity, ease of implementation, and monitoring requirements

- Simplicity
- Robustness
- Maximizing throughput for production lab use
- Minimizing sample transfers, extractions, filter steps, chemical additions (e.g., pH adjustments)

Validated Test Method 8327: PFAS Using External Standard Calibration and MRM LC-MS/MS





Non-Drinking Water Sample Methods: SW-846 Method 8327—Direct Injection

24 PFAS (including all target analytes in EPA Method 537)

Commercially available standards ("native" and isotopically labeled)

Direct injection-EPA Region 5/Chicago Regional Lab SOP

- Similar to draft American Society for Testing and Materials (ASTM) Method D7979
- Multi-laboratory validation study completed in 2018
- OLEM addressing public comments (closed August 26, 2019)
- Expected to be finalized in Fall 2020?

Target Quantitation Limits: 10 nanogram/L

Associated preparation method 3512 for aqueous matrices





Non-Drinking Water Sample Methods: CWA/SW846 Method—Isotope Dilution

More complex method relative to direct injection; however will:

- likely be more robust for complex matrices (e.g., wastewater influents, biosolids). Account for matrix effects (e.g., sorption) through isotopically marked standard recoveries;
- afford options to meet DoD requirements; and
- allow users to perform a deeper dive based on screening (e.g. 8327) results.

40 PFAS analytes-includes all analytes listed in 537.1, 533, and SW846 8327

Non-drinking water samples

- Surface water, groundwater, wastewater
- Landfill leachates
- Solids (soils, sediments, biosolids, tissues)



Non-Drinking Water Sample Methods: CWA/SW846 Method—*Isotope Dilution*

Build in flexibility

Columns
Elution schemes

Single laboratory validation in progress

- Collaborative effort among DoD, EPA Office of Water, EPA Office of Land and Emergency Management, and EPA ORD
- Assuming single lab validation success (target Dec. 2020), multi-laboratory validation will follow over 2021
- Method being developed in accordance with CWA and SWA-846 protocols for method development

Target Quantitation Limits: 1-10 nanogram/L

Matrices include:

- Wastewater (influent and effluent)
- Groundwater
- Surface water

- Landfill leachate
- Soil
 - Sediment
- Biosolids
 - Fish tissue



PFAS Analysis in Marine Waters

No EPA Approved method for PFAS in Marine Waters – ORD has a research method...

• LC-MS/MS Isotope dilution method

Method Details

- Covers 24 PFAS, Commercially available standards ("native" and isotopically labeled)
 - Similar to those in EPA 537.1, 533, and SW 846 8327
 - Target quantitation limits <1ng/L, with extracted samples
 - SPE sample concentration matrix elimination (up to 500 mL, Weak-Anion Exchange (WAX))
 - Accounts for matrix effects (e.g., high ionic strength and DOM)
- Adapted to estuarine sediments and TOP assay

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PFAS Analysis in Fish Tissue

No EPA Approved method for PFAS in fish tissue – EPA uses commercial laboratories' proprietary methods

- LC-MS/MS with solid phase extraction and isotope dilution
- Similar to DW 533 and SWA 1600 but don't dare call it a Modified Method...

Method Details

- Covers 13 carboxylic and sulfonic acids from C4 to C12, plus PFOSA; now 33 analytes
- Quantitation limits ranged from 0.25 to 1.25 ng/g (ppb) for the 13 (0.38-4.09 ng/g for 33)
- Spike stable isotopically labeled PFAS analogs into 1-2 g fillet tissue sample
- Sample digested with caustic (KOH or NaOH) methanol solution to release PFAS from tissue
- Solids removed by centrifugation, aqueous solution processed by SPE extraction
- Injected into LC-MS/MS for analysis

Longer chain PFAS C8+ most consistently present





Total Organofluorine Analysis using Combustion Ion Chromatography (TOF)





- No specialized or costly instrumentation required
- Applied for aqueous matrices and blood samples
- Removing the background inorganic F⁻ from the sample is important to make sure that the reported F⁻ is organic
- Can be developed on a wide commercial scale
- High priority for EPA. ORD working with OW-OST to develop a draft method in 2020

Set EPA

Total Organic Precursors (TOP)

- Developed by Houtz et al. No multi-laboratory validated standard methods
- Available from some contract laboratories
- Does not identify individual precursor compounds

Heated oxidative conversion

60 mM Persulfate, 125 mM Base

Oxidizable PFAA Precursors

85 °C for 6 Hrs

Total Oxidizable Precursors = PFCAs after oxidation – PFCAs before oxidation

- Applicable for aqueous and solid matrices
- Conservative estimate of the total concentration of PFAA precursors
- More expensive; sample needs to be analyzed twice for PFAS

PFCAs



Summary: EPA PFAS Methods, July 2020

EPA has validated Standard Methods complete or in development for PFAS in water

- Final SDWA Methods 533 and 537.1 for available for drinking water (29 PFAS)
- Method SW846-8327 validated and undergoing final review for non-potable water (24 PFAS)
- Method CWA-1600 undergoing single and multi lab validation for non-potable water and solids (40 PFAS)

EPA has or is developing additional methods for partner use

- Fish Tissue Isotope dilution method for 13 PFAS has been used in national surveys
- Serum Isotope dilution method (targeted and non-targeted) used in biomonitoring
- Ambient air and emissions Sampling and analysis methods undergoing development and testing
- Total Organic Precursors (TOP) Identify total PFAS load which may degrade to most persistent PFAS
- Total Organic Fluorine (TOF) Potential rapid screening tool to identify total PFAS presence/absence
- High resolution mass spectrometry Continued development and application of HRMS methods for discovery of novel PFAS, suspect screening, and non-targeted analysis



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