

Office of Research and Development

# **Update on PFAS Analytical Methods**

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# Per- & Polyfluoroalkyl Substances (PFAS)



**€EPA** 

- A class of man-made chemicals
  - Chains of carbon (C) atoms surrounded by fluorine (F) atoms, with different terminal ends
  - Complicated chemistry thousands of different variations exist in commerce
  - Widely used in industrial processes and in consumer products
  - Some PFAS are known to be PBT:
    - Persistent in the environment
    - Bioaccumulative in organisms
    - Toxic at relatively low (ppt) levels

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# **Methods-Sampling**

## **Guidance to avoid cross contamination in sampling**

- No teflon
- Avoid contact with clothes, materials containing PFAS (e.g. some food wrappers)
- See:
  - Sampling guidance from MI <u>https://www.michigan.gov/documents/pfasresponse/General\_PFAS\_Sampling\_Guidance\_634597\_7.pdf</u>
  - Interstate Technology and Regulatory Council Fact Sheet: Site Characterization Considerations, Sampling Precautions, and Laboratory Analytical Methods for PFAS pfas-1.itrcweb.org/wp-content/uploads/2018/03/pfas fact sheet site characterization 3 15 18.pdf
- PFAS Quality Assurance Plan and Data Review issues <u>epa.gov/fedfac/technical-fact-sheet-perfluorooctane-sulfonate-pfos-and-perfluorooctanoic-acid-pfoa-0</u>



# **Targeted vs Non-Targeted**

**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 semiquantitate 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



# **Non-Targeted Analysis**

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



- 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**



# **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; QAed 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



# **Types of Standard Methods**

## **Three broad categories of EPA Standard Methods:**

- Safe Drinking Water Act Methods
  - https://www.epa.gov/dwanalyticalmethods
- Clean Water Act Methods
  - https://www.epa.gov/cwa-methods
- SW846 Methods
  - https://www.epa.gov/hw-sw846/guidance-methods-development-andmethods-validation-resource-conservation-and-recovery-act

These are generally targeted methods for solids and water



## **Drinking Water Method 537: Revision I**

- Update: External lab validation for additional analyt
  - Perfluoro-2-propoxypropanoic acid (GenX chemical HFPO-D
  - Potassium 9-chlorohexadecafluoro-3-oxanone-1-sulfonate (
  - Potassium 11-chloroeicosafluoro-3-oxaundecane-1-sulfonate
  - Sodium dodecafluoro-3H-4,8-dioxanonate (ADONA, CAS 958)
- Incorporated clarifications issued in EPA Technical Advisory epa. 09/documents/pfoa-technical-advisory.pdf
- Final published method (November, 2018) epa.gov/water-resear ۲ methods
- LC/MS/MS with internal standards. Single lab lowest concentratic lacksquarerange from 0.53-6.3 ng/L

	-CPA Document #: EPA/600/R-18/352				
METHOD 537.1 DETERMINATION OF SEL POLYFLUORINATED AL WATER BY SOLID PHAS CHROMATOGRAPHY/T (LC/MS/MS)	DETERMINATION OF SELEC SUBSTANCES IN DRINKING Y LIQUID CHROMATOGRAPHY/ I. <u>SCOPE AND APPLICATION</u> 1.1 This is a solid phase extraction spectromety (Lasse extraction polyfluorinated alkyl substance data have been generated in reag in the table below.	METHOD 537.1 TED PER- AND POI VATER BV SOLID P TANDEM MASS SPE (SPE) liquid chromato thed for the determinati so (PFAS) in drinking w sent water and drinking	VFLUORINATED ALKYL HASE EXTRACTION AND (CTROMETRY (LC/MS/MS) graphy/tandem mass on of selected per- and ater. Accuracy and processor		
7	Analyte* Hexafluoropropylene oxide dimer acid N-ethyl perfluorooctanesulforamid	Acronym HFPO-DA	water for the compounds listed Chemical Abstract Services Registry Number (CASDA)		
J.A. Shoemaker and D.R. Tettenhors	N-methyl perfluorooctanesulfonamidoacetic acid Perfluorobutanesulfonic acid Perfluorodecanoic acid	NEtFOSAA NMeFOSAA PFRS	13252-13-6 <sup>b</sup> 2991-50-6 2355-31-9		
J.A. Shoemaker (Office of Research and Development), B.K. Boutin (N	Perfluorododecanoic acid Perfluoroheptanoic acid Perfluorohexanesulfonic acid	PFDA PFDoA PFHpA	375-73-5 335-76-2 307-55-1		
	Perfluorohexanoic acid Perfluorononanoic acid Perfluorooctanesulfonic acid	PFH <sub>X</sub> S PFH <sub>X</sub> A	375-85-9 355-46-4 307-24-4		
	Perfluorooctanoic acid Perfluorotetradecanoic acid Perfluorotridecanoic acid	PFOS PFOA	375-95-1 1763-23-1 335-67 1		
	Perfluoroundecanoic acid 11-chlorocicosafluoro-3-oxaundecane-1-sulfonic acid 9-chlorohexadecafluoro 2	PFTA PFTrDA PFUnA	376-06-7 72629-94-8 2058-94-8		
NATION OFFI U.S.F	4.8-dioxa-3H-perfluorononanoic acid * Some PFAS are commercially available as ammonium, sodiur of the analytes as anions while the counterior is interaction.	9CI-PF3OUdS 9CI-PF3ONS ADONA	763051-92-9° 756426-58-1d 919005-14-4°		
* HEPO A more section of a second section is inconsequential. Analyses and polassium salts. Takis method measures all forms of the GenX processing and technology. * HEO-LPF3OLdS is available in the GenX processing and technology. * OCL-PF3OLdS is available in salt form (e.g. CASRN of Polassime salt is 83329-89.9). * ADONA is available as the sodium salt (on Case. CASRN of Polassime salt is 83329-89.9).					
<ol> <li>Minimum Reporting Level (MRL) is the lowest analyte concentration that meets Data Quality Objectives (DQOs) that are developed based on the intended use of this</li> </ol>					



# Drinking Water Method 537: "Modified"

- Method 537 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.





# **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
  - Il support the fifth Unregulated Contaminant Monitoring Rule
  - Released December 2019
    - https://www.epa.gov/dwanalyticalmethods/analytical-methods-developed-epa-analysis-unregulated-contaminants

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# **Drinking Water Method 533**

Method 533 <sub>C</sub>	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) <sup>3</sup>	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)	

# **\$EPA**

### 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)

https://www.epa.gov/hw-sw846/validated-test-method-8327and-polyfluoroalkyl-substances-pfas-using-external-standard





### 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)
- Finalize in March (?) 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.

#### 33 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, multi-laboratory validation will follow over Summer 2020
- Method being developed in accordance with CWA and SWA-846 protocols for method development

#### Target Quantitation Limits: 1-10 nanogram/L

Sediment

•

#### Matrices include:

- Wastewater (influent Landfill leachate Biosolids and effluent) • Soil • Fish tissue
- Groundwater
- Surface water



### **PFAS Analysis 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

Contact: David Katz & Mark Cantwell EPA/ORD/CEMM (katz.david@epa.gov; cantwell.mark@epa.gov)

# **SEPA**

## **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





**PFAS** Analysis in serum

#### No EPA Approved method for PFAS in serum – ORD has a research method...

- HRMS analysis protein crash (removal) and isotope dilution
- Kotlarz et al., 2020 method details (*in review EHP*)

#### **Method Details**

- Covers 10 legacy PFAS, and 10 perflouro ether carboxylic and sulfonic acids
- Target quantitation limits 0.1 to 2.1 ng/mL, with extracted calibration curves (calf serum)
- Good agreement (<12% CV) with PFOA, PFOS, PFNA and PFHxS in NIST SRM 1957 human serum
- Does not include SPE step
- 50 uL of serum followed by formic acid denature and acetonitrile protein crash with centrifugation
- Injected into LC-Orbitrap for analysis; can be modified for LC-MS/MS

Contact: Mark Strynar, EPA/ORD/CEMM (Strynar.mark@epa.gov)

**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



- 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



## Total Organofluorine Analysis using Combustion Ion Chromatography (TOF)

Adsorption of PFAS on to activated carbon / other sorbents





- No specialized or costly instrumentation required
- Applied for aqueous matrices and blood samples
- Removing the background inorganic F<sup>-</sup> from the sample is important to make sure that the reported F<sup>-</sup> 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



## **PFAS Source Emissions Measurements**

#### **No EPA Approved methods for PFAS Emissions – EPA developing OTM-45**

- Considering both sampling and analysis methods, targeted and non-targeted
- Diverse sources chemical manufacturers, commercial applications, thermal treatment incineration processes
- Methods needed for source characterization and for control technology evaluation

#### **Method Development Details**

- Semi/Non-Volatiles Performance-based, Modified Method 5 (MM5 see <a href="https://www.epa.gov/hw-sw846/sw-846-test-method-0010-modified-method-5-sampling-train">https://www.epa.gov/hw-sw846/sw-846-test-method-0010-modified-method-5-sampling-train</a>) train approach using isotope dilution, GC/MS targeted and non-targeted analysis
- Volatiles modified TO-15 using SUMMA canisters, GC/MS targeted and non-targeted analysis (see <a href="https://www3.epa.gov/ttnamti1/airtox.html">https://www3.epa.gov/ttnamti1/airtox.html</a> for methods)
- Surrogate Indicators measure PFAS as a class, e.g., Total Organic Fluorine (TOF)

#### Methods in development through state, industry collaborations



### **PFAS** Ambient Air Measurements

#### No EPA Approved methods for PFAS ambient air methods

- Considering both sampling and analysis methods, targeted and non-targeted
- Applications include fenceline monitoring for fugitive emissions, deposition, receptor exposure

#### **Method Development Details**

- Ambient/Near-Source Field deployable Time of Flight/Chemical Ionization Mass Spectrometer for real time detection/measurement
- Semivolatile PFAS Performance Based following guidance in EPA TO-13a,
- Volatile PFAS SUMMA canisters, sorbent traps, GC/MS targeted and non-targeted analysis



#### Methods in development through state, industry collaborations

Contact: John Offenberg, US EPA/ORD/CEMM (offenberg.john@epa.gov) 22



## Summary: EPA PFAS Methods, January 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 and solids (24 PFAS)
- Method CWA-1600 undergoing single and multi lab validation for non-potable water and solids (24 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

# **Sepa**

## For More Information

#### ORD Executive Lead for PFAS R&D

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