Demonstration of Low PFAS Background Associated with the TurboVap[®] LV Automated Solvent Evaporation System

Environmental laboratories require a system which is clean, reliable, and affordable to evaporate the extracts resulting from the preparation of samples for PFAS testing. This document will outline a protocol that was developed for the TurboVap[®] LV system to evaporate extracts in methanol to dryness consistent with typical environmental PFAS procedures. It will also show data resulting from this protocol which demonstrates the cleanliness of the system for PFAS compounds using a sample analogue.



Experimental Design

To simulate a standard PFAS extract which would result from either an automated or a manual solid-phase extraction (SPE) procedure, 15 mL of methanol was added directly to polypropylene centrifuge tubes (VWR p/n 21008-670). The centrifuge tubes were loaded onto the 48-position Multi Rack within a TurboVap[®] LV and the protocol provided in Table 2 was used to concentrate the samples to dryness. The dry extracts were reconstituted in 1-mL storage vials using 990 μ L of 96% methanol/4% water and 10 μ L of an internal standard mixture was added. The resulting extracts were analyzed on an LC-MS/MS for the target compounds given in Table 1.

Note: For more in-depth information on the preparation procedure or the materials and equipment used, please refer to

application note AN958 entitled "Manual Extraction of PFAS in Drinking Water in Compliance with EPA Method 537.1."

 Table 2. TurboVap* LV Concentration Protocol.

Bath Temp	60 °C
Evaporation Mode	Method (Ramp Gradient)
Manifold Setup	48 positions
Rack Row Height	120 mm*
Step 1	2.5 L/min for 15 min.
Step 2	3.0 L/min for 15 min.
Step 3	3.5 L/min for 45 min.

*The nozzle position was adjusted such that it was as far to the right as possible to give the user a clear view of the vortex within the tube.

Table 1. PFAS analytes examined as possible contaminants within the evaporation and analysis steps of a standard extraction procedure.

Target Analyte	Acronym	CASRN
Perfluorobutanesulfonic acid	PFBS	375-73-5
Perfluorohexanoic acid	PFHxA	307-24-4
Hexafluoropropylene oxide dimer acid	HFPO-DA	13252-13-6b
Perfluorohexanesulfonic acid	PFHxS	355-46-4
4,8-dioxa-3H-perfluorononanoic acid	ADONA	919005-14-4e
Perfluoroheptanoic acid	PFHpA	375-85-9
Perfluorooctanoic acid	PFOA	335-67-1
Perfluorooctanesulfonic acid	PFOS	1763-23-1
Perfluorononanoic acid	PFNA	375-95-1
9-chlorohexadecafluoro-3-oxanone-1-sulfonic acid	9CI-PF3ONS	756426-58-1d
Perfluorodecanoic acid	PFDA	335-76-2
Perfluoroundecanoic acid	PFUnA	2058-94-8
N-ethyl perfluorooctanesulfonamidoacetic acid	NEtFOSAA	2991-50-6
11-chloroeicosafluoro-3-oxaundecane-1-sulfonic acid	11CI-PF3OUdS	763051-92-9c
N-methyl perfluorooctanesulfonamidoacetic acid	NMeFOSAA	2355-31-9
Perfluorododecanoic acid	PFDoA	307-55-1
Perfluorotridecanoic acid	PFTrDA	72629-94-8
Perfluorotetradecanoic acid	PFTA	376-06-7



Results

The data shown in Table 3 was generated using a protocol with a Minimum Reporting Level (MRL) of 2 ng/L for all compounds. In the cases where certain compounds were available only in their salt form, the masses of the compounds were adjusted accordingly to correct for the salt content (Table 3). As typical of many environmental methods, for a system to be deemed "clean," it must demonstrate that any background present is less than one-third of the MRL (1/3 MRL). An examination of the background data collected shows that the highest observed concentrations detected for all analytes are approximately 3.5x lower than the 1/3 MRL limit given by typical environmental methods.

Table 3	Results	of PFAS	Evaporation	Background	Study	(recoveries	in ng/L).	

Deallasta	MRL Conc.	TurboVap [®] LV						
Replicate		1	2	3	4	5	6	
PFBS*	1.77	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	
PFHxA	2.0	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	
HFPO-DA	2.0	N.D.	N.D.	N.D.	0.002	0.006	N.D.	
PFHxS*	1.89	N.D.	N.D.	N.D.	N.D.	0.061	N.D.	
ADONA*	1.89	N.D.	N.D.	N.D.	N.D.	N.D.	0.004	
PFHpA	2.0	N.D.	0.030	0.012	0.031	0.024	0.012	
PFOA	2.0	0.073	0.069	0.050	0.071	0.071	0.063	
PFOS*	1.91	0.143	0.143	0.115	0.145	0.130	0.137	
PFNA	2.0	N.D.	0.015	0.013	0.005	0.014	N.D.	
9CI-PF3ONS*	1.86	N.D.	0.004	N.D.	0.005	0.002	0.003	
PFDA	2.0	N.D.	N.D.	N.D.	0.005	0.013	N.D.	
PFUnA	2.0	N.D.	N.D.	N.D.	0.026	N.D.	0.012	
NEtFOSAA	2.0	N.D.	N.D.	N.D.	N.D.	N.D.	N.D.	
11CI-PF3OUdS*	1.88	N.D.	0.006	N.D.	N.D.	0.003	N.D.	
NMeFOSAA	2.0	N.D.	N.D.	N.D.	N.D.	N.D.	0.013	
PFDoA	2.0	N.D.	0.004	N.D.	0.007	0.008	0.007	
PFTrDA	2.0	N.D.	0.003	N.D.	0.013	0.014	N.D.	
PFTA	2.0	N.D.	N.D.	N.D.	N.D.	0.004	N.D.	

*Analytes were used in their salt form and calculated concentrations were corrected to compensate where needed.

Note: Where "N.D." is indicative of the inability of the target peak to be separated from the system background.



Figure 1. PFAS Background of the TurboVap^{*} LV.

*Analytes were used in salt form and their calculated concentrations were corrected to compensate.



Discussion

Separating the sample preparation process into its major components and examining them independently allows for the systematic elimination of factors contributing to the background of the overall process. In this document, the extraction process for a typical PFAS environmental sample was eliminated, and the evaporation and analytical processes were examined using a sample analogue. The experiment demonstrates that the TurboVap[®] LV, along with the other supplies and equipment used, fulfill the requirements for PFAS background levels and are thus acceptable to use in the general production of PFAS data for environmental compliance work.

Ordering Information:

Part Number	Description
415000	TurboVap [®] LV Automated Solvent Evaporation System
414964	TurboVap [®] LV Multi Rack (48 Positions, 10-20 mm Tubes)

EUROPE

Main Office: +46 18 565900 Fax: +46 18 591922 Order Tel: +46 18 565710 Order Fax: +46 18 565705 order@biotage.com Support Tel: +46 18 56 59 11 Support Fax: +46 18 56 57 11 eu-1-pointsupport@biotage.com

NORTH & LATIN AMERICA

Main Office: +1 704 654 4900 Toll Free: +1 800 446 4752 Fax: +1 704 654 4917 Order Tel: +1 800 446 4752 Order Fax: +1 704 654 4917 ordermailbox@biotage.com Support Tel: +1 800 446 4752 us-1-pointsupport@biotage.com

JAPAN

Tel: +81 3 5627 3123 Fax: +81 3 5627 3121 jp_order@biotage.com jp-1-pointsupport@biotage.com

CHINA Tel: +86 21 68162810 Fax: +86 21 68162829 cn_order@biotage.com

cn-1-pointsupport@biotage.com

Tel: +82 31 706 8500 Fax: +82 31 706 8510 korea info@biotage.com

KORFA

kr-1-pointsupport@biotage.com INDIA Tel: +91 11 45653772 india@biotage.com Distributors in other regions are listed on www.biotage.com

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