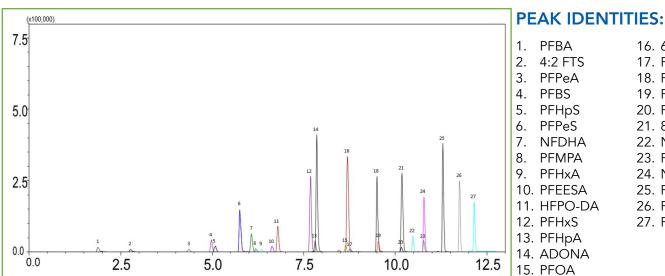
## HALO





## Analysis of 27 PFAS Compounds Using a Modified 1633 Method



## 16. 6:2 FTS 17. PFMBA 18. PFNA 19. PFOS 20. PFDA 21. 8:2 FTS 22. N-MeFOSAA 23. PFUnA 24. N-EtFOSAA 25. PFDoA 26. PFTrDA 27. PFTeDA

## **TEST CONDITIONS:**

Column: HALO 90 Å PFAS 2.1 x 100 mm, 2.7 µm Part Number: 92812-613 Delay Column: HALO® PFAS Delay 3.0 x 50 mm, 2.7µm Part Number: 92113-415 Mobile Phase A: 5 mM Ammonium Acetate Mobile Phase B: MeOH Gradient: %В Time 0.0 20 90 12.0 15.0 **0**0

15.0	70
15.1	20
18.0	END

Flow Rate: 0.4 mL/min Pressure: 489 bar Temperature: 44 °C Injection Volume: 1 µL Sample: LGC PFASiMix Product Number: DRE-A50000647MW Sample Concentration: 1µg/mL Sample Solvent: 96:4 Methanol/Water LC System: Shimadzu Nexera X2 ESI LCMS System: Shimadzu LCMS-8040 A mix of 27 PFAS standards was analyzed using a HALO® PFAS column. Due to the stability of PFAS compounds, there will always be a need for testing. Environmental agencies around the world have set strict limits to the amount of PFAS that can be found in drinking water. It is very important to quantify and identify these compounds in order to accurately determine whether the concentrations are within acceptable ranges.

By using the HALO® PFAS column, in conjunction with a modified EPA method, the 27 compounds above were separated with great resolution and peak shape in under 13 minutes. This method enables fast and robust PFAS separations.

MS Source Conditions: ESI -Spray Voltage: 4.5 kV Nebulizing gas: 2 L/min Drying gas: 17 L/min **DL temp:** 250 °C Heat Block: 400 °C





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