Sample preparation in Non-Target Screening for very polar compounds: Prospects and Difficulties

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Overview

1. The Challenge

2. The SPE Challenge

3. Applied Methods

4. Conclusion

1. The Challenge

Why to preconcentrate?

→ Many compounds can be in trace levels that even with Large Volume Injection are not detectable.

 \rightarrow Laboratories without "sensitive" instruments.

Until now:

Targeted analysis of polar compounds

• NORMAN screening trial of River Danube approach (Schymanski et al., 2015, Anal. Bioanal. Chem)

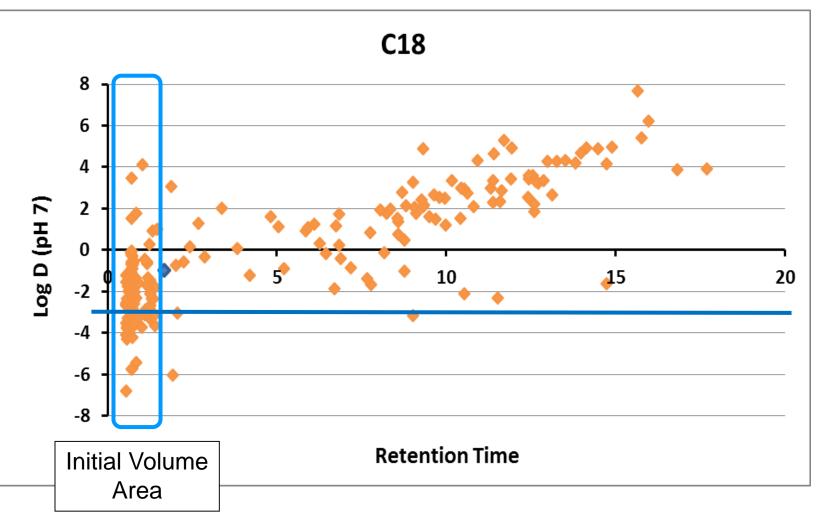
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Very Polar compounds ?

Compounds with LogD < -2.5 (pH7)

Not retained by:

- Polar embedded
- Polar Endcapped
- Non-Endcapped
 columns



Very Polar compounds ? Isocratic pump Binary pump 2 Binary pump 1 ESI-TOF-ZIC-HILIC Autosample-RP MS No HILIC **RP-C18** Λ HILIC **RP-C18**



2. The SPE Challenge



"How do you take the very polar compounds out of the most polar solvent: water?"



\rightarrow "Get rid" of the water first!

"Bumps" on our road!

\rightarrow The salt content of the sample

- Salt depositions trapping compounds.
- "Poisoning" of the [HILIC] absorbants.

 \rightarrow Time consuming and "harsh".



3. Applied Methods

Combination of RP SPE and HILIC SPE

 Lyophilisation (Freeze drying)
 Evaporation under Vacuum – BUCHI Syncore system

>{4. Membrane Nanofiltration}





4. Conclusion

Very polar compounds are becoming of increasing importance.

Sample preparation procedure should also adjust, if needed.

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THANK YOU!

What is your opinion on very polars?

