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Background

Per- and Polyfluoroalkyl Substances (PFAS) are chemicals used for many domestic and industrial purposes related to their physicochemical properties. However, those same properties make them mobile and persistent in the environment, and on top of that, they are toxic and can affect human health in the short and long term, as they are bio-accumulative. Many processes govern the transport of PFAS in the surface waters and groundwater, e.g. sorption, biodegradation, co-transport, and transformation. Monitoring PFAS at different locations can help understand these processes and provide datasets to calibrate and validate reactive transport models simulating PFAS fate and transport. This study compares PFAS presence and distribution in river water and groundwater at four Danube river sites. The site in Vienna has a clogging layer on top of the river bed and steady water levels while in Budapest, water levels are dynamic with no clogging.



Figure I: Study sites map

Methods

- Samples from the river and groundwater were collected bimonthly over one year in Vienna and Budapest (Surany, Tahi and Ráckeve). In Ráckeve composite sample from 3 production wells (PW) were taken.
- The analysis targeted 32 PFAS compounds and LCMS was used for PFAS analysis and pharmaceuticals.
- Censored data (<LOQ) were processed using regression on order statistics.
- Shapiro-Wilk test for normality and Kruskal-Wallis test for differences were performed on the data set.

Occurrence and Distribution of PFAS in the River and Groundwater at Two Danube Sites

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- ADONA had the highest concentration among all PFAS (average 10 ng/l), while many of the substances had analytical results below LOQ (Figure 3).
- * Background water affects PFAS concentrations in the production wells at Budapest sites.
- Some PFAS are possibly impacted from older water or different sources (i.e. PFOS, PFOA).
- Carbamazepine is behaving similar to PFAS while Diclofenac shows sorption to some extent.



Figure 3: Boxplots of PFAS concentrations in the Danube and bank filtered groundwater (boxes: 25 – 75%, line: median, whiskers: ±1.5 IQR%).

Figure 2: Piper diagram for Surany and Tahi

- Danube (p>0.05, Table 1).



- Majority of the PFAS concentrations are not normally distributed.
- Test of differences shows that there are no differences be sampling locations (p>0.05, Tab
- In Budapest, PFAS concentration Danube and groundwate (p<0.05).

Conclusion and future steps

- sorption or degradation for many of the compounds.
- coefficients.
- PhreeqC.

Figure 5: planned column setup (left), preliminary groundwater flow model for Vienna site (right).

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In Vienna, PFAS concentrations in all monitoring wells are similar as the

In Surany and Tahi, there is an influence from background water (Fig. 4).

Figure 4: PFAS concentration time series in the Danube and groundwater at Tahi. Table I: Kruskal-Wallis test P_values

rations are	PFAS	P_value			
		Surany	Tahi	Rackeve	Vienna
	PFBA	0.089	0.819	0.086	0.380
t in Vienna	PFHxA	0.045	0.606	0.014	0.602
atwoon all	PFHpA	0.512	0.963	0.327	0.430
etween all	PFOA	0.001	0.023	0.142	0.253
ble I).	GenX	0.011	0.030	0.624	0.705
ions in the	PFOS	0.001	0.030	0.221	0.918
ions in the	PFBS	0.296	0.918	0.462	0.930
er differ	PFHxS	0.000	0.022	0.027	0.547
	NaADONA	0.013	0.797	0.086	0.161
	PFPeA	0.053	0.984	0.027	0.382

PFAS are generally persistent during soil passage: there was no/minimal

PFOS and PFOA seem to be from older water at the Budapest transect.

Soil column experiments will provide more information about sorption

Modelling of PFAS transport using MODFLOW coupled with MT3DMS and





