Year: 2018

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DOI: https://doi.org/10.1021/acs.est.8b02763

Trends in Micropollutant Biotransformation along a Solids Retention Time Gradient

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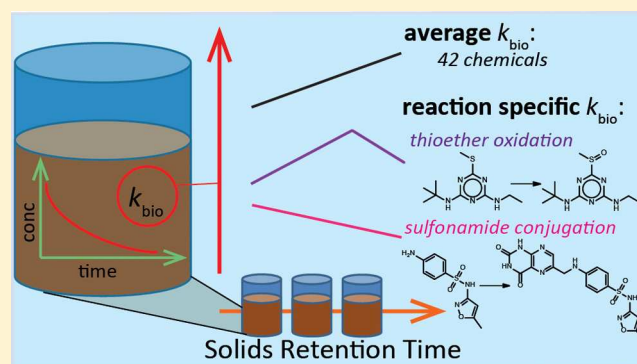
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S Supporting Information

ABSTRACT: For many polar organic micropollutants, biotransformation by activated sludge microorganisms is a major removal process during wastewater treatment. However, our current understanding of how wastewater treatment operations influence microbial communities and their micropollutant biotransformation potential is limited, leaving major parts of observed variability in biotransformation rates across treatment facilities unexplained. Here, we present biotransformation rate constants for 42 micropollutants belonging to different chemical classes along a gradient of solids retention time (SRT). The geometric mean of biomass-normalized first-order rate constants shows a clear increase between 3 and 15 d SRT by 160% and 87%, respectively, in two experiments. However, individual micropollutants show a variety of trends. Rate constants of oxidative biotransformation reactions mostly increased with SRT. Yet, nitrifying activity could be excluded as primary driver. For substances undergoing other than oxidative reactions, i.e., mostly substitution-type reactions, more diverse dependencies on SRT were observed. Most remarkably, characteristic trends were observed for groups of substances undergoing similar types of initial transformation reaction, suggesting that shared enzymes or enzyme systems that are conjointly regulated catalyze biotransformation reactions within such groups. These findings open up opportunities for correlating rate constants with measures of enzyme abundance such as genes or gene products, which in turn should help to identify enzymes associated with the respective biotransformation reactions.

INTRODUCTION

To assess and minimize the risks imposed by the release of anthropogenic chemicals into the environment, understanding which factors determine transport and degradation processes of chemicals at the end of their product life-cycle is important. Pharmaceuticals and other domestic and industrial chemicals are frequently detected in wastewater.[1] Although activated sludge-based treatments are primarily optimized to reduce biological oxygen demand (BOD) and nutrients such as ammonium and phosphorus, they also remove micropollutants (MPs) to varying degrees. In fact, for many polar, organic micropollutants, biotransformation by activated sludge microorganisms is an important removal process.[1] Numerous attempts have been made to understand the influence of the molecular structure, process-related parameters, properties of the wastewater or characteristics of the microbial communities on MP biotransformation.[2−8] Although previous work sheds light on the importance

of various factors, a significant part of the observed variability in MP removal under different wastewater treatment conditions remains unexplained.[9,10] Yet, a more thorough understanding of how observed variabilities in MP biotransformation rates and pathways, including formation of potentially hazardous transformation products (TP), depend on different influencing factors is of significant interest: It would enable more accurate predictions of biotransformation, which in turn would support chemical risk assessment, and it would open up opportunities to optimize treatment processes toward improved micropollutant removal.

Solids retention time (SRT), or sludge age, which defines the average time microbial biomass resides in a given treatment

unit, has repeatedly been discussed as having a major influence on micropollutant biotransformation. In several studies, positive relationships between SRT and micropollutant removal have been reported[1,11−26] and a beneficial effect of SRT on MP removal is widely accepted.[27] Clara et al.[12] first reported a strong correlation between effluent concentrations and SRT for bisphenol A, bezafibrate, ibuprofen and natural estrogens. Ternes and Joss observed that nitrifying wastewater treatment plants (WWTPs) operated at SRTs above 10 d removed MPs more efficiently than WWTPs operated at less than 4 d SRT.[1] However, in other studies, contrasting results were reported of no connection between MP removal and SRT,[10,19,28−33] or even higher removal at lower SRTs for certain MPs.[28]

To further understand these contradictory findings, there is a need to distinguish between overall removal of MPs and biomass-normalized (or biomass activity-normalized) biotransformation rate constants. As elaborated in Schwarzenbach et al.,[34] pseudo-first-order kinetics have frequently been observed for MP biotransformation in microbial communities, allowing to calculate biomass-normalized rate constants (often denoted k_{bio}).[1,35−37] At fixed reactor volumes and hydraulic retention times (HRTs), an increase in SRT leads to a higher concentration of active biomass.[15] Even if biomass-normalized biotransformation rate constants were not affected by SRT, those higher biomass concentrations would most likely instigate a higher percentage of biological removal.[22,38,39] A number of more recent studies describe attempts to better understand the influence of SRT on biotransformation rate constants. Petrie et al. have shown that estrogen biotransformation per bacterial cell was higher at 10 d than at 3 d SRT.[38] Falås et al. found higher VSS-normalized biotransformation rate constants for ketoprofen and naproxen in activated sludge from WWTPs operated at an SRT of at least 7 d compared to low SRTs of 1−3 d.[23] In contrast, others reported that a decrease in SRT from 20 to 5 d led to higher k_{bio} values for diclofenac, ibuprofen, naproxen and caffeine.[32] Similarly, in a comparison of three activated sludge systems operated at SRTs of 3, 10, and 20 d, the endocrine disrupting substances bisphenol A, triclosan and 4-n-nonylphenol showed highest k_{bio} values at 3 d of SRT.[19] Finally, no systematic influence of SRT on k_{bio} for different MPs was observed for activated sludge reactors operated in parallel at high SRTs of 25, 40, and 80 d.[33]

Taken together, previous observations remain restricted to a limited number of substances and do not allow for any general conclusions on the influence of SRT on MP biotransformation rates. It remains unclear which substance classes and types of biotransformation reactions are positively associated with SRT and through which mechanisms SRT affects biotransformation rates and pathways. Therefore, in this study, we aimed to address the question how MP biotransformation depends on SRT using MPs from different substance classes undergoing various types of biotransformation reactions. Specifically, we wanted to explore whether consistent trends would be found for classes of substances undergoing similar biotransformation reactions, and whether these trends could serve to generate hypotheses about underlying mechanisms. For this purpose, we established six bioreactors operated at a gradient of SRTs and designed a strategy to consistently profile the biotransformation potential of the cultivated activated sludge communities. The strategy included the selection of a diverse and sufficiently large set of MPs that were expected to undergo a range of oxidation and substitution reactions, and the establishment of

an efficient workflow to study the biotransformation rate constants and reactions of the selected MPs.

■ METHODS

Reactor Setup and Operation. For cultivation of activated sludge at different SRTs, six 12 L reactors were operated in parallel. The reactors were connected to a programmable logical controller (Wago 750-881) and a SCADA system (Citect V7.2, Schneider Electric), enabling automated operation as previously described.[33] Online sensors allowed to control fill levels (Cerebar PMC131, Endress+Hauser) and dissolved oxygen (DO) concentrations (Oxymax COS61D, Endress+Hauser) and to monitor temperature (ISEmax CAS40D, Endress+Hauser). The reactors were equipped with feed pumps, discharge valves, flow-controlled fine-bubble aerators and stirrers. For inoculation of the reactors, activated sludge was collected from the nitrification tank of a full-scale WWTP receiving mostly municipal wastewater (WWTP Niederglatt, 40 000 population equivalents, details in the Supporting Information (SI), section S1). DO levels in the reactors were controlled within the range of 1.5−3 mg/L via intermittent aeration. The six reactors were operated as sequencing batch reactors (SBRs) at a fixed HRT (12 h) and different SRTs (1, 3, 5, 7, 10, and 15 d). The reactors were operated in fully automated 4 h cycles comprising a settling phase, a discharge phase, a feed phase and a reaction phase as detailed in the SI (section S1). For reactor feeding, municipal wastewater was collected after primary treatment (screening, grit removal, and sedimentation) from an urban catchment of 30 000 population equivalents. To adjust the SRTs, different portions of activated sludge were withdrawn from the six reactors according to Table S1. All total suspended solids (TSS) measurements performed prior to and during the biotransformation experiments showed gradually increasing values from the first to the sixth reactor, confirming a gradual increase in actual SRTs (SI section S2). The reactor temperatures varied slightly according to the temperature in the sewer (18 ± 2 °C).

Biotransformation Profiling Strategy and Workflow. Our strategy to profile the biotransformation potential of the activated sludge communities is summarized in the following. More details are given in the SI, sections S2−S7.

Selection of Chemicals. Multiple substances covering the same chemical class and for which we expected similar initial biotransformation reactions according to molecular structure-based *in silico* pathway prediction using the EAWAG pathway prediction system (EAWAG-PPS, http://eawag-bbd.ethz.ch/predict/) and reported pathways from literature were included. For instance, for primary amides, nitriles or carboxylic esters, hydrolysis reactions were expected, whereas oxidative demethylation reactions were expected for phenylureas or generally for tertiary amides. Chemicals of known environmental relevance were additionally included, resulting in a total of 93 analyzed chemicals (for further details on chemicals and expected transformation reactions, see section S3).

Biotransformation Experiments. Biotransformation batch experiments were started after 48 days (Exp1) and 187 days (Exp2) of reactor operation with a total of 77 (Exp1) and 93 (Exp2) chemicals, respectively. In Exp1, the experiment was conducted in the 12 L SBRs directly where, for the duration of the batch experiment, the cycling was paused after a feed phase. In Exp2, the activated sludge was transferred to smaller glass bottles (100 mL, triplicates for each SRT) that were equipped with caps with two holes to maintain aerobic conditions and

placed on a shaker table. MPs were spiked to final concentrations of 6 μg/L each in each reactor. Samples were collected at multiple time points over 3−4 days. The samples were centrifuged, and the supernatant was transferred to HPLC vials. A mix of isotope-labeled internal standard substances was added to account for losses and interferences during liquid chromatography coupled to high-resolution mass spectrometry (LC-HRMS) measurements, and the samples were stored at 4 °C for a maximum of 7 days until analysis. In the first 4 h of each experiment, samples were collected for measuring concentrations of NH_4^+, NO_2^- and NO_3^-. pH was measured using a HQ30d Flexi Meter (Hach Lange). In parallel to each biotransformation experiment, sorption and abiotic control experiments were performed in triplicate in 100 mL reactors according to Gulde et al.,[37] i.e., autoclaved filtrate (cellulose nitrate filter, 0.45 μm, Sartorius Stedim Biotech) and autoclaved activated sludge were used to estimate the fractions that were abiotically degraded and sorbed to activated sludge solids, respectively.

Additional experiments to investigate biotransformation under oxygen limited conditions (ExpOx) and to support transformation product analysis (ExpTP) were started after 356 days and after 243 days of reactor operation, respectively, both with activated sludge sampled from the reactor operated at 7 d of SRT. In ExpOx, biotransformation under aerobic conditions (as in Exp1 and Exp2) was compared to biotransformation in sealed bottles run under DO limitation. In these, the initially present electron-accepting nitrogen species (nitrite and nitrate) were consumed in the first 5 h, and anaerobic conditions then prevailed until the end of the experiment. For further details on biotransformation experiments see section S4.

Chemical Analysis and Evaluation of Rate Constants. Mass spectra were recorded on a QExactive Plus (Thermo Scientific) mass spectrometer. Full-scan MS spectra were acquired in both positive and negative ionization modes and the acquisition of MS[2] fragmentation spectra was triggered at m/z values corresponding to masses of suspected transformation products (see below). Details on chromatographic separation and mass spectrometric analysis are provided in the SI (section S5). Calibration standards were prepared in nanopure water (Barnstead Nanopure, Thermo Scientific) covering a concentration range between 0.05 and 10 μg/L.

Target chemical concentrations were quantified with Tracefinder 3.1 (Thermo Scientific). From the resulting concentration−time series, first-order biotransformation rate constants were derived in line with the pseudo-first-order assumption, which has proven useful to describe biotransformation of low concentrated chemicals in cases where the enzymatic turnover of the substrate is rate-limiting (for further discussion, see Section S6).[1,34] First-order rate constants were calculated by a censored linear regression applied to the logarithmic concentrations against time, using the software R (Version: 3.3.0) and the command "censReg" from the package "censReg". Concentrations below 5% of the initially spiked concentration (for which for some MPs no reasonable chromatographic peak, i.e., less than three full-scans, and deviation from first-order kinetics was observed) were treated as censored values. Chemicals selected for further analysis included those that (1) did not strongly adsorb (<30% in the sorption experiments), (2) did not show abiotic degradation (<20% at end of the abiotic control experiments) and (3) exhibited first-order kinetics (see Section S6 and Table S6, also for criteria for first-order kinetics). To study the effect of differences in microbial community composition across reactors containing different total biomass concentrations, first-order rate constants for these chemicals were converted into second-order rate constants by normalization with TSS. In doing so, uncertainties of the observed rate constants and TSS measurements were propagated using a Monte Carlo approach (R package "propagate"). Biases involved with using TSS as an approximation for active biomass are discussed in the SI (section S2). Because adsorption to activated sludge and abiotic losses were minor for the selected chemicals, their normalized rate constants are further denoted as second-order biotransformation rate constants, k_{bio}. Finally, to enable comparisons of rate constants across all reactors, only chemicals were further considered for which k_{bio} for at least one SRT was higher than a predefined threshold value (Exp1, 0.162 L g_{TSS}^{-1} d^{-1}; Exp2, 0.121 L g_{TSS}^{-1} d^{-1}) corresponding to approximately 30% removal over 2 days in the reactor with the lowest measured biomass concentration.

Trends of biotransformation rate constants with SRT were analyzed as follows: We calculated Spearman rank correlation coefficients (ρ) between k_{bio} and SRT by repeated ($n = 10\,000$) sampling of k_{bio} values for each SRT from their distributions obtained from error propagation. We further compared trends with SRT across MPs by substance-wise autoscaling and centering of k_{bio} values (details on scaling in section S6).[40] Heatmaps were produced using the R package "pheatmap" and ordering of chemicals followed hierarchical clustering (using Euclidean distance and complete linkages). For ExpOx, k_{bio} values were obtained similarly as for Exp1 and Exp2 (see Table S10), and a ratio between rate constants under anaerobic and aerobic conditions, further denoted k_{anaer}/k_{aer}, was calculated for each chemical.

Analysis of Transformation Products and Assignment of Reaction Types. A suspect transformation product screening was performed using the software Compound Discoverer 2.0 (Thermo Scientific). After peak picking and prefiltering (details in section S7), a subset of features was selected based on a comparison with a predefined suspect list. In addition to the TPs predicted by the Eawag-PPS or described in the literature, further potential TP masses were calculated considering mass shifts of common biotransformation reactions (hydroxylation, dihydroxylation, demethylation, dehydrogenation, hydrogenation and decarboxylation). To facilitate structure elucidation and assessment of the parent-TP relationships, an auxiliary experiment (ExpTP) was performed in which the micropollutants were spiked in groups of 10−20 chemicals and at higher concentrations (50 μg/L each). If for detected features with exact masses contained in the suspect list a reasonable peak area-time trend (either increasing or first in- and then decreasing) was observed in MS spectra from Exp2 and the TP was only detected in samples from batches spiked with the respective parent MP in ExpTP, MS[2] fragmentation spectra were analyzed (further details in section S7) unless the TP could be confirmed by comparing the retention time with an authentic reference standard (Table S11).

TP evidence, i.e., observed changes in molecular formula and structure, was then used to assign corresponding reaction types to the parent MPs. This approach involved varying degrees of uncertainty because for most TPs their relative importance could not be accurately quantified because of a lack of authentic standards. Also, despite having assembled a rather exhaustive TP suspect list, TPs might still have been missed in our analysis. However, in most cases, the TPs found were in good agreement with previously published pathways. Additionally, based on ratios of k_{anaer}/k_{aer} observed for MPs with well-established
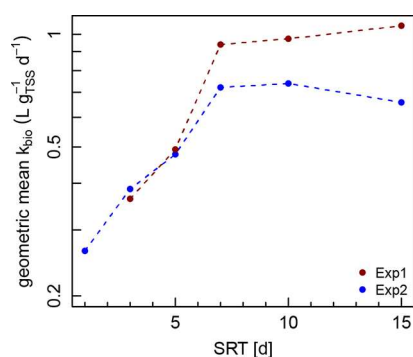
Figure 1. Geometric mean rate constants (logarithmic y-axis) obtained from Exp1 (33 MPs) and Exp2 (42 MPs) in dependence on the SRT.

transformation reactions (e.g., pargyline, valsartan and atenolol), we could confirm our expectation that k_{anaer}/k_{aer} ratios would generally be smaller for oxidative transformations than for substitution-type reactions. These ratios were therefore used to support assignment of reaction types to MPs with less well-established transformation reactions.

■ RESULTS AND DISCUSSION

Reactor Operation. After inoculation, the six parallel sequencing batch reactors were operated at SRTs of 1, 3, 5, 7,

10 and 15 d as described in the Methods section. During the initial phase, TSS as well as influent and effluent concentrations of nitrogen species (NH_4^+, NO_2^-, NO_3^-) were measured on a regular basis (Tables S2 and S3). After 7 days of operation, TSS values had developed from initially 3.4 g/L in all reactors to 0.2, 1.3, 1.7, 2.0, 2.9 and 3.1 g/L for SRTs of 1, 3, 5, 7, 10 and 15 d, respectively. The reactor at 1 d SRT had lost its nitrifying activity after 7 days while the reactors at 7, 10, and 15 d SRT remained nitrifying at all measured time points. Possibly caused by fluctuations in temperature,[41] conversion of ammonium to nitrate was also observed at 3 and 5 d of SRT in Exp2. Indeed, slightly higher mean temperatures were recorded in the 2 weeks before Exp2 (19.3 ± 1.0) compared to Exp1 (18.1 ± 0.7). Data on TSS, pH, temperature, nitrifying activity and oxygen uptake rates during Exp1 and Exp2 are provided in Tables S4 and S5.

Dependence of Biotransformation Rate Constants on SRT. Of the initially selected MPs, a considerable number (Exp1, 20; Exp2, 16) was not degraded sufficiently to allow for a reliable determination and comparison of k_{bio}. In particular, the biotransformation of MPs with a triazine structure (for which halide substitution was predicted) and MPs predicted to undergo O-dealkylation or tertiary amide N-dealkylation were often only slowly transformed unless an alternative (faster) transformation pathway was possible (Table S6, for observed

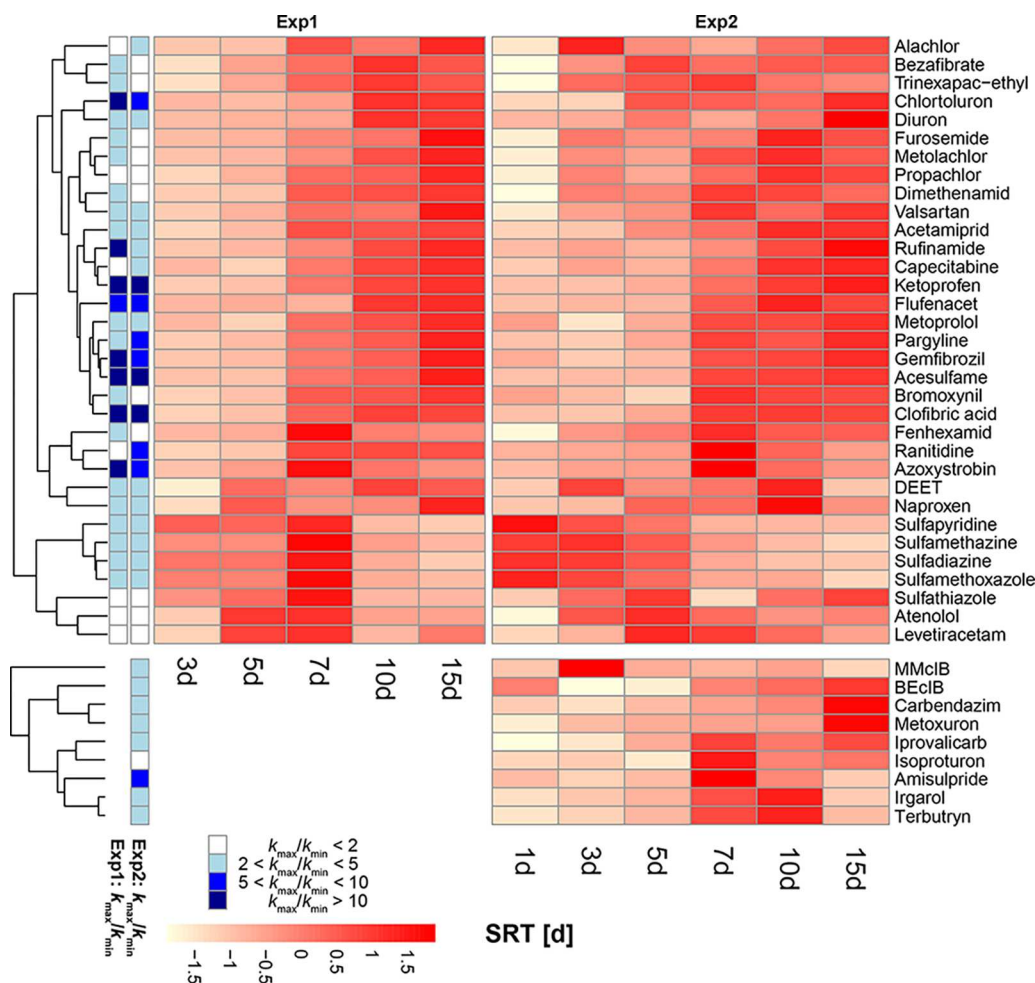**Figure 2.** Substance-wise autoscaled rate constants from Exp1 (33 MPs) and Exp2 (42 MPs) (see SI section S6 for details). Levels of relative differences in k_{bio} between 15 and 3 d of SRT are annotated on the left. Ordering of MPs followed hierarchical clustering of the combined trends obtained in Exp1 and Exp2 (33 MPs) and, separately, for the 9 MPs additionally investigated in Exp2.

Table 1. Reactions Observed for Each Substance and Additional Experimental Information

reaction type	name	observed reaction[a]	Exp1: k_{max}/k_{min}[b]	Exp2: k_{max}/k_{min}[b]	ρ (Exp1)[c]	ρ (Exp2)[c]	k_{anaer}/k_{aer}[d]	k_{inh}/k_{cntrl} (ATU)[e]	k_{inh}/k_{cntrl} (OCT)[e]
oxidation	Chlortoluron	dealk./dihydrox./hydrox.	>3.36	6.48	0.9 (0.8–1.0)	0.6 (0.6–0.9)	<0.08	0.16	0.73
	Diuron	dealk.[43]/dihydrox.	3.20	3.34	0.9 (0.6–1.0)	0.7 (0.5–0.9)	<0.14	0.27	0.60
	Isoproturon	dealk.[10]/hydrox.	n/a	1.75	n/a	0.6 (−0.1–0.7)	<0.13	0.55	0.58
	Metoxuron	dealk./dihydrox.	n/a	2.74	n/a	0.9 (0.2–1.0)	0.36	<0.80	<0.80
	Irgarol	S-oxidation[44]	n/a	3.36	n/a	0.0 (−0.1–0.7)	<0.16	0.45	0.68
	Ranitidine	S-oxidation[f,45] N-oxidation[45]	1.65	6.00	0.7 (0.5–1.0)	0.6 (0.3–0.6)	0.32	0.22	0.70
	Terbutryn	S-oxidation[44]	n/a	4.82	n/a	0.4 (0.0–0.7)	<0.13	0.38	0.74
	Valsartan	amide dealk.[36]	3.69	2.11	0.9 (0.7–1.0)	0.8 (0.6–0.9)	0.04	0.90	0.64
	BEclB	amide dealk./hydrox.[36]	n/a	2.50	n/a	1.0 (0.8–1.0)	<0.14	0.77	0.43
	MMclB	amide dealk.[36]	n/a	2.68	n/a	−0.7 (−1.0 − −0.2)	<0.35	0.72	<0.25
	Furosemid	amine dealk.	4.78	1.72	1.0 (0.9–1.0)	0.6 (0.2–0.9)	<0.10	0.14	0.54
	Amisulpride	N-oxidation/amine dealk.	n/a	7.99	n/a	0.3 (−0.2–0.6)	<0.34	0.51	0.98
	Pargyline	N-oxidation[42]	3.54	5.38	0.9 (0.6–1.0)	0.9 (0.6–1.0)	0.05	0.29	0.55
	Ketoprofen	hydrox./biphenyl pathway[46]	19.47	17.06	1.0 (0.9–1.0)	1.0 (1.0–1.0)	<0.01	0.71	0.89
	Gemfibrozil	hydrox.[47]	14.21	6.60	1.0 (0.9–1.0)	1.0 (0.8–1.0)	<0.01	0.94	0.74
	Clofibric acid	hydrox.	>11.60	10.17	0.9 (0.9–1.0)	0.7 (0.6–1.0)	0.08	0.33	0.50
	Iprovalicarb	hydrox.	n/a	3.15	n/a	0.7 (0.7–0.9)	0.18	0.79	0.93
both	Capecitabine	hydrox./hydrolysis	1.75	2.02	0.9 (0.8–0.9)	0.9 (0.7–1.0)	0.09	0.33	0.30
	Bezafibrate	amide dealk./hydrolysis[48]	3.05	1.62	0.9 (0.7–1.0)	0.4 (0.1–0.8)	0.03	0.49	0.52
substitution	Levetiracetam	amide hydrolysis[48]	1.37	1.98	0.1 (−0.6–0.7)	0.0 (−0.4–0.1)	0.06	0.70	0.71
	Atenolol	amide hydrolysis[10]	1.66	1.66	0.1 (0.0–0.2)	−0.8 (−0.9 − −0.2)	1.97	1.21	1.09
	Rufinamide	amide hydrolysis	15.84	4.54	1.0 (1.0–1.0)	0.9 (0.9–0.9)	0.37	0.65	0.87
	Dimethenamid	substitution/reduction[g]	2.07	1.21	1.0 (0.8–1.0)	0.5 (−0.5–0.9)	0.70	0.88	0.91
	Alachlor	substitution/reduction[g]	1.65	2.23	0.9 (0.8–0.9)	−0.1 (−0.2–0.3)	0.67	0.95	0.98
	Flufenacet	substitution/reduction[g]	5.96	5.38	0.9 (0.8–0.9)	0.8 (0.8–0.9)	0.72	0.77	0.81
	Propachlor	substitution[10]/reduction[g]	1.99	1.54	1.0 (0.9–1.0)	0.8 (0.2–1.0)	0.92	0.90	0.88
	Metolachlor	substitution/reduction[g]/dealk.	2.32	1.28	1.0 (0.9–1.0)	0.6 (−0.3–1.0)	0.75	0.89	0.87
	Sulfamethazine	pterin-conjugation[49]	2.99	2.57	−0.6 (−0.7 − −0.3)	−1.0 (−1.0 − −0.9)	0.43	1.09	1.05
	Sulfapyridine	pterin-conjugation[49]	3.83	3.16	−0.7 (−0.7 − −0.6)	−1.0 (−1.0 − −0.7)	0.63	1.18	1.04
	Sulfadiazine	pterin-conjugation[49]	2.57	2.26	−0.6 (−0.7 − −0.6)	−0.9 (−1.0 − −0.7)	0.65	1.11	1.03
	Sulfamethoxazole	pterin-conjugation[49]	2.89	2.65	−0.7 (−0.7 − −0.5)	−0.9 (−1.0 − −0.8)	0.89	1.20	1.10
	Sulfathiazole	pterin-conjugation[49]	1.56	1.28	−0.5 (−0.7–0.2)	0.1 (−0.7–0.8)	0.61	0.74	0.88
	Azoxystrobin	ester hydrolysis[10]	22.91	6.74	0.6 (0.6–0.6)	0.6 (0.1–0.6)	0.38	0.91	0.84
	Trinexapac-ethyl	ester hydrolysis[10]	2.27	1.38	0.9 (0.7–0.9)	−0.5 (−0.9–0.3)	0.36	0.80	1.07
	Bromoxynil	nitrile hydration	3.61	1.52	1.0 (0.8–1.0)	0.6 (0.2–0.9)	0.62	0.62	0.92
	Acetamiprid	nitrile hydration	3.77	3.43	0.9 (0.7–1.0)	0.9 (0.9–1.0)	0.24	0.50	0.39
	Carbendazim	carbamate hydrolysis[45]	n/a	3.30	n/a	1.0 (0.8–1.0)	0.19	<0.48	0.84
unknown	Acesulfame	n/a[h]	74.02	23.67	1.0 (0.9–1.0)	0.9 (0.6–1.0)	0.08	0.97	1.08
	Metoprolol	n/a[i]	2.32	2.67	0.9 (0.6–0.9)	0.9 (0.6–1.0)	<0.02	n/a	n/a
	DEET	n/a[h]	2.48	2.47	0.8 (0.4–0.9)	−0.3 (−0.6 − −0.1)	0.02	0.92	0.97
	Naproxen	n/a[h]	2.53	3.96	0.7 (0.3–0.7)	0.3 (0.3–0.4)	0.02	1.03	1.04
	Fenhexamid	n/a[h]	3.06	1.66	0.6 (0.3–0.7)	0.6 (0.3–0.7)	0.28	0.41	0.66

Table 1. continued

[a]Observed reactions (for details, see methods and SI, section S7). Dealk. indicates dealkylation, hydrox. indicates hydroxylation, dihydrox. indicates dihydroxylation. Citations refer to literature describing the same transformation in activated sludge or activated sludge-derived communities. [b]Ratios of highest rate constants over lowest rate constants per substance within the range of SRTs between 3 and 15 d. For chlortoluron and clofibric acid in Exp1, the minimum rate constants were very low and set to 0.048 L/(g_{TSS} × d), corresponding to 10% removal over 2 days in the reactor operated at 1 d SRT. [c]Spearman rank correlation coefficients and empirical confidence intervals (95%). [d]Ratio of rate constants obtained under aerobic and anaerobic conditions (rate constants in Table S10). [e]Ratio of rate constants (inhibited over control) using the ammonia monooxygenase inhibitors allylthiourea (ATU) and octyne (OCT). Data based on Men et al.,[50] rate constants are provided in Table S10. [f]Quantification of N- and S-oxides indicated that S-oxidation represents the major reaction. [g]For the acetanilides, reductive dehalogenation was observed besides the substitution reactions. [h]For acesulfame, DEET, naproxen and fenhexamid, no TP could be confirmed in the suspect TP screening. [i]Metoprolol has been described to form atenolol acid in activated sludge.[45] Because metoprolol was not spiked and was, therefore, present in lower concentrations than atenolol, potential minor contribution to the formation of atenolol acid from metoprolol remained elusive.

pathways see below). For a number of MPs, concentration–time curves deviated from an exponential decay insofar as the transformation seemed to stall after an initial decline in concentration. In agreement with the recently described trapping of amine-containing MPs in protozoa,[42] this pattern was mainly observed for amine-containing MPs.

For the 33 (Exp1) and 42 (Exp2) substances selected for further analysis according to the criteria defined above, biomass-normalized rate constants k_{bio} were calculated (Figure S3 and S4). The comparably low biomass concentration in the reactor at 1 d SRT (0.43 ± 0.03 g/L) in Exp1 led to low measured first-order rate constants with high relative errors (Figure S5). Therefore, this reactor was not considered for further analyses. For both experiments, geometric mean k_{bio} values across all compounds were calculated for each SRT (Figure 1). An increasing trend of the mean k_{bio} with SRT was observed between SRTs of 3 and 7 d (Exp1, +160%; Exp2, +87%), whereas mean k_{bio} values at 7, 10 and 15 d SRT were rather similar. Considering that fractions of active biomass per total suspended solids are expected to decrease with higher SRT[15,35] (also see section S2), normalization by TSS likely led to a slight underestimation of the overall increase in biotransformation efficiency per active cell with SRT.

When looking at trends of individual MPs instead of mean rate constant trends, we observed a remarkable variety of differing trends. In Figure 2, the scaled rate constants for the individual MPs in both Exp1 and Exp2 are presented as a clustered heatmap. Additionally, the heatmap is annotated with information on the ratio k_{max}/k_{min} between the lowest and the highest rate constant observed between 3 and 15 d SRT. k_{max}/k_{min} ratios close to 1 indicate that the microbial communities were similarly efficient in biotransformation of the respective substance at different SRTs, whereas high k_{max}/k_{min} ratios indicate that changes in the microbial communities along the SRT gradient strongly affected the respective transformation.

The trends shown in Figure 2 confirm that the majority of micropollutants feature increasing degradation with increasing SRTs. However, for individual substances, different trends can be observed. Although some MPs roughly follow the trend of the geometric mean presented in Figure 1, many showed highest rate constants only at the highest SRTs of 10 or 15 d. Also, opposite trends with lowest rate constants at highest SRTs (e.g., for the sulfonamides) or highest rate constants at intermediate SRTs (e.g., for ranitidine, azoxystrobin, DEET or naproxen) were observed. Yet, the generally similar trends across Exp1 and Exp2 demonstrate that biotransformation rate constants of various MPs show systematic trends with SRT and that these trends are reproducible at different time points after start of the reactor operation and across different experimental

settings (12 L bioreactors in Exp1 versus 100 mL bioreactors in Exp2).

Substance Class- and Reaction Type-Specific Trends. The diversity of observed trends suggests that the different microbial species and enzymes responsible for the biotransformation of individual substances depend differently on SRT. Therefore, we hypothesized that clustering of trends could potentially be indicative of interactions of a specific (group of) enzyme(s) with similar functional moieties of different MPs. To investigate this hypothesis, we compared trends within and among different classes of substances and reaction types as characterized based on observed transformation products and additional information, e.g., k_{anaer}/k_{aer} (Table 1).

Sulfonamides. A consistent trend across all five sulfonamide antibiotics (sulfamethazine, sulfapyridine, sulfadiazine, sulfamethoxazole and sulfathiazole) was confirmed by hierarchical clustering (Figure 2, Table 1). With a slight, but continuous decrease along the SRT gradient in both Exp1 and Exp2, overlaid with a maximal k_{bio} at 7 d SRT in Exp1, the sulfonamides show a dependence on SRT that is distinctly different from all other substances. For all five investigated sulfonamides, TPs related to the recently described pterin-sulfonamide conjugation pathway were predominantly detected.[49] This transformation pathway, presumably initiated by the enzyme dihydropteroate synthase (DHPS), is related to the folate synthesis pathway that is ubiquitously present in bacteria and linked to bacterial growth. Because the TSS-normalized heterotrophic activity is expected to be slightly decreasing with higher SRTs due to the presence of more slow-growing organisms,[15,35] this hypothesis is consistent with the negative trend of both sulfonamide biotransformation rate constants and TSS-normalized oxygen uptake rates (OUR) with SRT (see Tables S4 and S5). The reason for the increased OUR and biotransformation rate constants in Exp1 at 7 d SRT is not known. As expected, neither inhibition by ATU or OCT nor oxygen limited conditions affected sulfonamide biotransformation. The previously reported sulfamethoxazole cleavage product 3-amino-5-methylisoxazole was not detected.[51]

Acetanilides. The acetanilides alachlor, dimethenamid, flufenacet, metolachlor and propachlor generally showed consistent, increasing trends with SRT. Yet, except for flufenacet (k_{max}/k_{min} > 5), which holds an oxyacetanilide moiety rather than a chloroacetanilide moiety, the acetanilides generally showed relatively small differences across the reactors (k_{max}/k_{min} 1.28−2.32). From soil degradation studies, the major known biotransformation pathway for acetanilides is initial glutathione-S-transferase (GST)-mediated conjugation followed by further transformation to the corresponding oxanilic acid (OXA) and ethanesulfonic acid (ESA) transformation products.[52] In this study, TPs related to this pathway were

detected for all investigated acetanilides. The GST system is a defense system ubiquitously present in plants, bacteria and animals. Our observation of only slight dependence of k_{bio} on SRT thus seems plausible given the expectation that, even if the microbial community composition changes with SRT, most microbes present should be able to catalyze this transformation. Additionally, reductive dehalogenations and, for metolachlor, an oxidative demethylation were observed (Table 1). Consistent with the suggested dominance of the GST-mediated biotransformation pathway, all five acetanilides showed little dependence of k_{bio} on oxygen limited conditions (k_{anaer}/k_{aer}: 0.67−0.92) or inhibition by ATU or OCT (k_{inh}/k_{cntrl} > 0.7).

Nitriles. For bromoxynil and acetamiprid, transformation of the nitrile to the corresponding amide was confirmed in the TP analysis, whereas the suspected further reaction of the amide to the carboxylic acid was not detected. Both MPs showed comparable increasing trends and a positive correlation of their rate constants with SRT (Figure 2, Table 1). Whereas nitrilases typically catalyze the reaction of nitriles to carboxylic acids, biotransformation by nitrile hydratase enzymes would explain the formation of the observed amides.[53,54] Unlike bromoxynil, acetamiprid showed inhibition by ATU and OCT and a stronger dependence on aerobic conditions, which could indicate that an additional, presumably oxidative pathway (not detected) was present.

Esters. As observed earlier,[10] the two esters trinexepac-ethyl and azoxystrobin showed different patterns although carboxylic acid TPs, suggesting an ester hydrolysis reaction, were detected for both MPs (Table 1). Most recently, it was demonstrated that, in activated sludge, azoxystrobin seemed to be nearly exclusively transformed by protozoan hydrolases[55] and protozoa composition is indeed expected to depend on SRT.[56,57] In our study, a strongly increased rate constant for azoxystrobin in the 7 d SRT reactors was observed in Exp1 and Exp2 (Figure 2). In contrast, no strong trends along SRT was observed for trinexapac-ethyl, which is consistent with the fact that no effect of protozoa inhibition was detected for trinexapac-ethyl,[55] and suggests that its hydrolysis is catalyzed by other enzymes, e.g., esterases, that are widely present in bacteria.[58]

Primary and Secondary Amides. For the three primary amides atenolol, levetiracetam and rufinamide, formation of the corresponding carboxylic acid, presumably via a hydrolysis reaction, was observed. Whereas the aliphatic amides atenolol and levetiracetam showed a very similar trend and low k_{max}/k_{min} ratios (<2) across SRTs, rufinamide, which contains an aromatic, more sterically hindered primary amide group, showed a strong positive correlation with SRT (ρ, 0.9−1.0; k_{max}/k_{min}, 4.5−15.8). Generally, amidohydrolases are widespread among bacteria,[58] supporting the observation for atenolol and levetiracetam that the reaction readily occurs at different SRTs. In contrast, rufinamide hydrolysis seems to rely on more specific enzymes that show a strongly increasing abundance with higher SRT.

As atenolol and levetiracetam, the secondary amides bezafibrate, fenhexamid and amisulpride did not reveal strong relationships with SRT. For bezafibrate, the detected TPs suggest that both amide hydrolysis and N-dealkylation occurred, whereas for fenhexamid the predicted hydrolysis was not observed. For amisulpride, TPs corresponding to N-oxidation and amine dealkylation were detected.

Tertiary Amides. For the tertiary amides valsartan, DEET, BEclB and MMclB low k_{anaer}/k_{aer} ratios (<0.35) suggest that mainly oxidative reactions are relevant. However, no consistent trends and reactions were observed. It has earlier been observed that tertiary amide dealkylation (except for valsartan) was relatively slow,[36] explaining why many tertiary amides were not biotransformed fast enough to allow for a comparison of k_{bio} here (Table S5). Similarly, this may explain why for the acetanilides, which are also tertiary amides, amide dealkylation was not the preferred pathway. Possibly, also for DEET and BEclB, other reactions might outcompete N-dealkylation, potentially explaining the different trend in k_{bio}.

Phenylureas. Chlortoluron and diuron were investigated in both Exp1 and Exp2 and are clustered together (Figure 2). Both showed an increase in k_{bio} with SRT, pronounced k_{max}/k_{min} values (3.2−6.5) and small values of k_{anaer}/k_{aer} ratios (<0.15). In Exp2, isoproturon and metoxuron were additionally investigated. For both MPs, k_{bio} increased with SRT and k_{anaer}/k_{aer} ratios of <0.13 (isoproturon) and 0.36 (metoxuron) were observed. As chlortoluron and diuron, metoxuron showed the highest k_{bio} at 15 d SRT. Demethylation at the urea-N was observed for all four phenylureas. Additionally, dihydroxylated transformation products were detected for diuron, chlortoluron and metoxuron and oxidation of a benzyl group to a carboxylic acid group was observed for isoproturon and chlortoluron. Likely, differences in observed trends are therefore caused by a different relative importance of the detected pathways.

Thioethers. The triazines irgarol and terbutryn (only investigated in Exp2) showed a similar trend with highest k_{bio} at 7 and 10 d SRT, strongly resembling the trend observed for ranitidine in Exp2 (Figure 2). For all three MPs, the S-oxide TP was the major metabolite detected as described previously for irgarol and terbutryn.[44] Additionally, the biotransformation of all three MPs is affected to a similar degree by inhibition with ATU (k_{inh}/k_{cntrl} 0.22−0.45), whereas only weak inhibition by OCT was found (k_{inh}/k_{cntrl} 0.68−0.74), and all k_{bio} values were considerably smaller under anaerobic conditions (k_{anaer}/k_{aer} < 0.33). The observation of this characteristic trend, together with the detected TPs, may therefore provide a hint that these three MPs are biotransformed by the same or related enzymes, most likely a specific type of a monooxygenase.

MPs with Particularly Strong Positive Dependence on SRT. For several MPs, high Spearman rank correlation coefficients were observed together with high k_{max}/k_{min} ratios (Table 1). The five MPs with both ρ > 0.9 and k_{max}/k_{min} > 10 in at least one of the two experiments are acesulfame, ketoprofen, clofibric acid, gemfibrozil, and rufinamide. The artificial sweetener acesulfame is present in high concentrations in wastewater and has previously been suggested as stable tracer substance.[59] However, more recently, it has been found that acesulfame was degraded by activated sludge communities from different WWTPs.[50,60,61] Castronovo et al. observed quantitative biotransformation of acesulfame to sulfamic acid and a metabolic degradation was hypothesized but the reaction pathway is not entirely understood yet.[60] We are not aware of any previous studies having demonstrated the observed dependence of acesulfame biotransformation on SRT. For ketoprofen, clofibric acid and gemfibrozil, higher removal percentages were observed in an MBR system operated at 80 d compared to 20 d SRT.[15] In the case of ketoprofen, higher biotransformation rate constants for nitrifying sludges (from WWTPs operated at higher SRTs) compared to non-nitrifying sludges were observed by Falås et al.[23] Since in their study and also here no strong inhibitory effect of ATU was observed, a causal relationship with nitrification again seems unlikely.

Overall, a rather weak or even negative dependence on SRT was observed for many substitution-type transformations. In contrast, most oxidative transformations displayed clear trends of increasing degradation with SRT and often high k_{max}/k_{min} ratios (Table 1, also see Figure S6 in the SI showing the relationship between $\log(k_{anaer}/k_{aer})$ and $\rho \times \log(k_{min}/k_{max})$). Also, at least three out of the five chemicals showing strong correlations with SRT underwent oxidative transformations. Since microbial community composition and function can be expected to change with SRT,[62] our findings suggest that the observed substitution reactions are less dependent on these changes than the oxidative transformation reactions observed. Indeed, many of the observed substitutions are plausibly catalyzed by common enzymes involved in central metabolism or general defense mechanisms (e.g., GST, DHPS, esterases, peptidases). In contrast, the observed oxidative transformations are mostly expected to be catalyzed by oxidoreductases such as monooxygenases, which are known to be rarer (i.e., less generally widespread among different bacterial species[58]) and highly differentially expressed. The strong dependence of oxidative biotransformation reactions on SRT thus suggests that certain enzymes like monooxygenases either become more abundant or diverse at higher SRTs.

Potential Mechanisms of SRT Dependence. Generally, mainly two hypotheses have previously been discussed for increasing biomass-normalized MP biotransformation rate constants with increasing SRT: (1) Higher SRTs might allow slow-growing bacteria to establish stable populations, leading to increased metabolic capacity at higher SRTs (e.g,[15,63]). (2) The changed nutrient availabilities (lower food to microorganisms ratio) at higher SRTs may cause changes in enzyme expression patterns (e.g,[15,16,62]). Also, both mechanisms have been hypothesized to lead to a higher taxonomic and/or functional diversity, which had been associated with increased MP removal earlier.[58,64]

With respect to hypothesis (1), one prominent explanation for higher MP biotransformation rates at higher SRTs is the increased presence of nitrifying microorganisms that are only able to grow above a certain threshold of SRT. Nitrifying bacteria, and, in particular, the enzyme ammonia monooxygenase (AMO), have previously been associated with micropollutant biotransformation.[10,65−69] In fact, in our study, the increase in nitrifying activity in the range between 1 and 7 d SRT (Tables S4 and S5) coincided with the strongest increase in mean MP biotransformation rates. However, the biotransformation of many MPs that exhibited increasing trends were not or only slightly affected by inhibitors of nitrifying activity (e.g., acesulfame, ketoprofen, gemfibrozil or rufinamide in Table 1), suggesting that AMO is likely not the primary driver of increased biotransformation rate constants at higher SRTs for the majority of substances studied. With respect to hypothesis (1) another interesting observation was that there were barely any obvious outliers in the trends (i.e., intermediate SRTs at which the k_{bio} values of individual substances showed distinct peaks, see Figures S3 and S4). If specific metabolic pathways of specific key species were responsible for certain transformation reactions, fluctuations in their abundance would translate into fluctuations in biotransformation rates. Therefore, in our study either the abundance of the involved key species is very stable and does, for many MPs, predominantly depend on the SRT, or, what we consider more likely, the biotransformation of most MPs is less dependent on individual species but is rather

achieved by a range of microorganisms and enzymes as hypothesized by de Lorenzo.[70]

Regarding hypothesis (2), in parallel to the observed positive trends for most oxidations, we observed decreasing trends for concentrations of effluent chemical oxygen demand (COD) toward higher SRTs (Table S3). Since higher cytochrome P450 enzyme expression has been observed under conditions with lower availability of easily degradable carbon,[71] and because many cytochrome P450 enzymes can transform a broad range of substrates, these observations support hypothesis (2).

A few MPs, however, showed exceptional patterns. The sweetener acesulfame was not only the MP with by far the highest influent concentrations (>50 μg/L), but it also showed extraordinarily high k_{max}/k_{min} ratios (Exp1, 74.0; Exp2, 23.7), supporting the hypothesis of a very specific (and possibly growth-related) pathway.[60,61] Specific pathways were also suggested for two more of the five substances showing strongest positive dependence on SRT, i.e., gemfibrozil[47] and ketoprofen.[46] Azoxystrobin was the only MP that showed high k_{max}/k_{min} ratios (Exp1, 22.9; Exp2, 6.7) but low correlation with SRT because of an outstanding maximum at intermediate SRTs. As discussed above, this MP might be transformed by specific protozoan species with a high abundance in the 7 d SRT reactor.

Implications. In this study, we demonstrated a clear increase in average biomass-normalized biotransformation rate constants with increasing SRT across an extensive list of MPs, independent of any effects caused by increased biomass concentrations or hydraulic retention times.[22] The fact that not only average rates but also many individual trends showed pronounced positive relationships revealed a clear direct influence of SRT on micropollutant biotransformation. If, in specific wastewaters, individual MPs whose biotransformation rates strongly benefit from high SRTs are of particular concern (e.g., for toxicological reasons or due to an industrial point source), operation at higher SRTs may thus provide a cost-effective solution to remove those MPs. Because we could further show that the strong observed increase from 3 to 10 d SRTs is not solely linked to the nitrifying activity, our results suggest that micropollutant removal may also benefit from increased SRTs in wastewater treatment situations in which nitrification is hindered, for instance under low temperature conditions[25,72] or high BOD loadings.[73] However, the fact that for a number of MPs (e.g., the class of sulfonamide antibiotics) their rate constants did not increase with increasing SRTs (or remained below detectable limits under all SRTs tested), supports previous reports stating that a complete removal of micropollutants in domestic wastewaters likely cannot be achieved with biological treatment solutions alone.[33]

While we noted a large diversity of different trends with SRT for individual compounds, emphasizing the general complexity of the relationship between SRT and biotransformation rate constants, remarkable similarities in biotransformation patterns were observed for MPs undergoing similar transformations (e.g., sulfonamides, thioethers, nitriles, phenylureas). This does not only imply that transformation product and pathway analysis is beneficial for the understanding of how operational parameters affect micropollutant removal but it may also help to obtain a more thorough understanding of biotransformation in complex microbial communities in general. Biotransformation of micropollutants present at low concentrations in mixed microbial communities is poorly understood at the level of involved enzymes and/or microorganisms. One promising approach to start elucidating these relationships may be to

mine for associations between rate constants and the abundance of enzymes, enzyme-encoding genes or gene transcripts.[74] Compared to results from pure culture experiments whose relevance to mixed communities has been questioned,[75] this approach allows to study biotransformation without strongly interfering with the system. Our results support the future application of such an approach in two ways: First, classification of substances according to their main transformation reaction type allows to specifically search for associations of rates with the abundance of enzymes (or the encoding genes or gene transcripts) that are known to catalyze the observed type of biotransformation reaction and to thus reduce false positive correlations.[74] Second, the fact that micropollutants with shared functional groups indeed showed similarities in observed reactions and trends supports the validity of the underlying hypothesis of association mining, i.e., that similar transformation reactions of different micropollutants are catalyzed by the same (group of) enzyme(s). Finally, if micropollutants with similar functional groups can indeed be expected to be biotransformed by the same enzyme systems, this should also facilitate prediction of transformation rates based on molecular structure and community functional information in the future.

■ ASSOCIATED CONTENT

Ⓢ Supporting Information

The Supporting Information is available free of charge on the ACS Publications website at DOI: 10.1021/acs.est.8b02763.

> Details on reactor operation; details on biotransformation experiments including details on mass spectrometric analysis; supplementary results from biotransformation experiments including rate constants obtained in Exp1, Exp2 and ExpOx; detailed results from transformation product analysis (PDF)

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Notes

The authors declare no competing financial interest.

■ ACKNOWLEDGMENTS

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