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No evidence of a human influence on the mineralization of terrestrial dissolved organic matter (DOM) in Alpine fluvial networks

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Abstract

The influence of human activities on the role of inland waters in the global carbon (C) cycle is poorly constrained. In this study, we investigated the impact of human land use on the degradation of terrestrial dissolved organic matter (DOM) and its potential impact on bacterial respiration in ten independent catchments of the Lake Geneva Basin. Sites were selected along a gradient of human disturbance (agriculture and urbanization), and we monitored bacterial respiration in parallel to DOM bioavailability. Fluorescence spectroscopy was used to determine the influence of human land uses on DOM sources and composition as well as the dynamic of degradation or production of the fluorophores identified in our study sites. Higher bacterial respiration measured in agro-urban streams related to a short-term bioreactive pool (0-6 days of incubation) from autochthonous origin, whose the size increased with human disturbance. On the other hand, the degradation of terrestrial DOM was not impacted by human activities and was not found to contribute substantially to aquatic bacterial respiration. Although human land uses controlled DOM sources, composition and bioavailability at the basin scale, our results showed that human activities in the Lake Geneva Basin had a limited impact on the net C exchanges between inland waters and the atmosphere related to DOM mineralization. Considering that greater proportion of autochthonous-DOM signature is a common feature in human-impacted catchments, the implications of our results likely apply more widely. Yet, on a global scale, the influence of human activities remains to be determine given the large diversity of effects of agriculture and urbanization on freshwater DOM depending on the local environmental context.

1. Introduction

Continental surface waters receive more terrestrial carbon (C) than they export toward oceans, leading to the conceptualization of inland waters as active pipes that process, emit and store C during their travel from lands to oceans (Cole et al., 2007). Within this framework,

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the mineralization of terrestrial dissolved organic matter (DOM) by aquatic heterotrophic bacterial communities is a key process by which terrestrial C returns to the atmosphere through CO_2 emissions (Fasching et al., 2014; Lapierre et al., 2013; Mayorga et al., 2005) and whose flux is significant at the global scale (Battin et al., 2008). The study of DOM transport and transformation along fluvial networks is therefore of primary importance, yet a major gap in knowledge revolves around the impact of human activities on the reactivity and bacterial use (i.e., respiration or allocation to biomass) of terrestrial DOM in inland waters (Creed et al., 2015; Xenopoulos et al., 2021).

Agricultural and urban land uses are major catchment features impacting DOM sources and composition in aquatic ecosystems from local (Wilson and Xenopoulos, 2009) to regional scales (Lambert et al., 2017; Williams et al., 2016). Streams draining agricultural landscapes and/or urban catchments are commonly enriched in DOM of low molecular weight compared to streams draining forested catchments whereby DOM is dominated by aromatic, high molecular weight compounds (Lambert et al., 2017; Williams et al., 2010). Greater proportion of lower molecular weight compounds in agro-urban streams can be the consequence of a greater autochthonous algal production and bacterial activity in nutrient-enriched waters (Fuß et al., 2017; Lu et al., 2014; Williams et al., 2010; Wu et al., 2019), reduced hydrological connection with terrestrial sources (Giling et al., 2014; Parr et al., 2015), or transfer of less humified soil organic matter due to agricultural practices (Humbert et al., 2020; Lambert et al., 2017; Landsman-Gerjoi et al., 2020). As lower molecular weight molecules are typically more labile and easily available for uptake to bacterial communities (Berggren et al., 2010; Catalán et al., 2017; Kaplan and Bott, 1989), greater DOM processing can be expected in agro-urban streams (Hosen et al., 2014; Parr et al., 2015). However, the destabilization of a stock of soil organic material built before the conversion of forests or wetlands for agriculture or urban development can lead to the mobilization of large amount of humic and highly aromatic DOM into surface waters (Ekblad and Bastviken, 2019; Graeber et al., 2012; Hu et al., 2016; Petrone et al., 2011). As a consequence, the impact of human land uses on the dynamic of DOM in inland waters may be highly diverse depending on how agriculture and urbanization affect DOM sources, content, and composition as well as external drivers such as inorganic nutrients known to regulate bacterial DOM processing (Guillemette and del Giorgio, 2012; Reche et al., 1998).

Different scenarios about the consequences on net C exchanges for surface waters can be envisaged depending on the impact of human land uses on freshwater DOM. First, increasing delivery of colored and aromatic terrestrial DOM can lead to increase CO_2 emissions supposing that this terrestrial material gets respired by bacterial communities (Fasching et al., 2014; Lapierre et al., 2013). Second, increase in the export of low molecular weight DOM, either derived from terrestrial sources or produced in-stream, can results in more

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DOM respired and emitted as CO_2 into the atmosphere (Bodmer et al., 2016; Borges et al., 2018). However, if changes in DOM composition result from an enhancement in aquatic primary production, enhanced respiration of autochthonously-produced DOM shall not lead to higher net CO_2 emissions since the amount of C emitted into the atmosphere would be lower or equivalent to the amount of CO_2 previously fixed by primary aquatic producers. An alternative scenario would be that the release of simple and labile organic compounds derived from autochthonous sources enhances the degradation of terrestrial aromatic DOM by the so-called priming effect (Bianchi, 2011), inducing a final net increase of C emitted to the atmosphere. The priming effect is a process through which labile pool of DOM can enhance ("prime") the degradation of a more recalcitrant DOM pool based on interactions between microbial communities and/or changes in their functions (Guenet et al., 2010; Kuzyakov et al., 2000) but its occurrence in aquatic ecosystems is highly debated (Attermeyer et al., 2014; Bengtsson et al., 2018; Lambert and Perga, 2019). To determine how human activities may impact the mineralization of terrestrial DOM and infer the consequence on the active role of inland waters into the global C cycle, it is therefore necessary to evaluate the consequences on DOM sources, composition, but also DOM bioavailability and to identify which fraction of the DOM pool fuels respiration. This information is critical to establish the role of human land uses on the linkage between terrestrial and aquatic ecosystems.

In this study, we aimed to investigate the impact of human land uses on the role of inland waters as bioreactors with regard to the processing of terrestrial DOM. Water samples were collected at two periods in ten independent catchments selected along a gradient of human pressure (agriculture and urbanization) in the Lake Geneva Basin. Patterns in DOM degradation were investigated based on standardized degradation experiments and by the consumption of specific compounds of low molecular weights. Decreases in dissolved organic carbon (DOC) concentrations and changes in DOM composition (assessed by fluorescence spectroscopy) during incubations were used to unravel the contribution of different fractions to the bulk DOM kinetic degradation as well as to the bacterial respiration measured under field conditions. With this approach combining field work and laboratory experiments, we specifically aimed to identify the origin of DOM contributing to bacterial respiration in human-impacted streams and to evaluate the impact of human activities on the mineralization of terrestrial DOM in Alpine fluvial networks.

2. Material and methods

2.1 Study sites and sampling

Stream and river waters were collected in ten independent tributaries of the Lake Geneva, the largest lake of Western Europe located at the border between France and Switzerland in the Western Alps (Figure 1). Lake Geneva is situated in the Alpine foreland

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between the Alps and the Jura mountains and was carved during Quaternary glaciations mostly into the Tertiary Molasse. Drainage areas of streams and rivers ranged from 11 to 5240 km^2, Strahler order from 2 to 7, mean elevation from 614 to 2124 m, and land cover was dominated by forests (39±19%), croplands (30±22%), grasslands (13±14%), and urban areas (11±14%) according to the Swiss Federal Office for the Environment (FOEN). Agriculture is dominated by non-irrigated arable lands and vineyards, and forests by coniferous and broad-leaved forests. Selected streams and rivers drained a mosaic of land cover categories and were classified as agro-urban or forest-grassland streams if the sum of croplands and urban areas extents was higher or lower than 50%, respectively (Table 1).

Samples were collected on two occasions, at the end of autumn between the 13th and 14th of November 2018 and at the end of winter between the 5th and 7th of March 2019. Water temperatures during field campaigns were 10.1±1.8 °C and 6.6±1.1 °C, respectively, but discharge and precipitation conditions at the time of collection were similar according to some data provided by the FOEN. Water was collected below surface in 2 L acid-washed high-density polyethylene (HDPE) bottles and filtered on site. A known volume of water (between 1000 and 1500 mL) was filtered on pre-combusted (450°C for 4 h) Whatman glass fiber filters (GF/F grade, 0.7 µm nominal pore size, 47 mm diameter) for chlorophyll a (Chl*a*) measurements. This filtered water was either stored in 1L acid-washed HDPE bottles for further use in the incubation experiments and the measurements of bacterial metabolism, as described below, or further filtered at 0.2 µm with polyethersulfone (PES) syringe encapsulated filters for dissolved organic carbon (DOC), colored and fluorescent DOM (CDOM and FDOM), soluble reactive phosphorus (SRP), and dissolved inorganic nitrogen (DIN) measurements. Syringe encapsulated filters were first rinsed with ultrapure water (60 mL) in the laboratory and then with 15-20 mL of water from the sampling site before collecting samples. Samples for DOC concentrations were stored in 40 mL acid-washed glass vials with polytetrafluoroethylene (PTFE)-coated septa, and samples for CDOM and FDOM were stored in 40 mL acid-washed amber glass vials with PTFE-coated septa. Samples for SRP, and DIN were stored separately in 50 mL sterile centrifuge tubes. All samples and filters were brought back to the laboratory within 3 hours in cool and dark conditions. Samples for SRP and Chl*a* measurements were frozen at -21°C until analysis. Other samples were stored in a dark chamber at 4-5°C until analysis typically done within the following two weeks.

2.2 Characterization of DOM degradation kinetics

Incubations were prepared once back to the laboratory. Water samples previously filtered on site at 0.7 µm were divided into 250 mL acid-washed glass flasks and incubated for 28 days in the dark at 20°C. Different subsets of flasks (in triplicates) were prepared and sacrificed for DOC measurements and DOM characterization every 3-5 days for the first 10

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days and then every 5-8 days up to day 28. Samples were filtered at 0.2 µm with PES syringe encapsulated filters as described above and stored in a dark chamber at 4-5°C. DOC measurements were done within 48h after collection, and CDOM and FDOM analyses over the following week. Dissolved oxygen depletion during the incubations was avoided by leaving a large headspace within the glass flasks and by a regular (every 3-4 days) renewal of the headspace.

Several models can be used to characterize DOM degradation kinetic. Here we applied a first-order exponential decay model in order to derive a decay constant describing the overall dynamic of DOM degradation (Lambert and Perga, 2019; Lu et al., 2013; Shang et al., 2018) as well as the size of the short- and a long-term reactive carbon pools (STRC and LTRC, respectively). Decreasing DOC concentrations during the incubations were modeled using GraphPad Prism 8 software according to the following equation:

$$DOC(t) = DOC_{cons} \times e^{-k*t} + DOC_{residual}$$

where DOC(t) is the DOC concentration measured at the incubation time t, DOC_{cons} the amount of DOC consumed at the end of the incubation, *k* the decay constant, and $DOC_{residual}$ the concentration of the residual pool remaining in solution at the end of the incubation. Biodegradable DOC (BDOC) was calculated as the difference in DOC between the initial and final time. Furthermore, we used the *k* decay constant from the model to quantify the STRC and LTRC pools: STRC was defined as the amount of DOC consumed within the first six days of incubation and the LTRC as the amount of DOC degraded between days 6 and 28. The separation between STRC and LTRC pools was based on a breakpoint in the degradation curves observed around the 6^{th} day of incubation in almost all experiments. Finally, changes in DOM composition during the incubations were also monitored by fluorescence measurement coupled to parallel factor analysis (PARAFAC) as described below.

2.3 Degradation of low molecular weights compounds

We also determined the consumption of low molecular weight compounds including carbohydrates (CAR), carboxylic and acetic acids (C&AA), and amino acids (AA) using Biolog Ecoplates ® (Garland and Mills, 1991; Weber and Legge, 2009, 2010). Ecoplates ® are 96-well microplates containing 31 different carbon substrates (in triplicates) plus a tetrazolium dye. The bacterial respiration activity associated with a specific substrate reduces the tetrazolium dye and produces a color measurable at 590 nm in absorbance. The intensity of color development in color can be related to the amount of substrate consumed (e.g., Berggren and del Giorgio 2015). Water from each site (125 µL) filtered at 0.7 µm was added to each well of one Ecoplates ® per site, which was then incubated in the dark at 20°C for 3 to 9 days until the average well color development (AWCD) reached an asymptote. The absorbance at 590 nm was measured with a TECAN microplate reader one to two times per day. Color

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development of each carbon substrate was calculated as the blank-corrected absorbance at the time when the AWCD was closest to 0.5 (0.51±0.11, n = 60) following recommendations of Weber and Legge (2010). Carbon substrates were then classified as CAR, C&AA, or AA according to Weber and Legge (2009). The color development for each of these categories was calculated as the mean of the color of the respective substrates normalized by the AWCD. While Ecoplates ® also include other organic compounds such as polymers or amines/amides, we focused in this study on the consumption of low molecular weight compounds known to support bacterial respiration (Kaplan and Bott, 1989).

2.4 Bacterial respiration measurements

The 0.7 µm-filtered waters used for incubation experiments were also used to measure bacterial respiration (BR) in stream and river waters, ensuring comparability between water quality and bacterial metabolism. BR was determined from the decrease in dissolved oxygen (DO) in 60 mL borosilicate serum bottles filled with water, sealed with a butyl stopper and crimped with an aluminum cap without headspace (3 serum bottles per site). These vials were equipped with SP-PSt7 oxygen planar sensor spots (PreSens) in order to follow DO consumption after 24h of incubation in the dark at 20°C. Initial (1h after the start of incubation) and final DO was recorded using a PreSens Fibox 4 equipped with a fiber-optic oxygen transmitter. Calibration of the PreSens Fibox 4 (two-point calibration at 0 and 100% oxygen saturation) was performed and verified before measurements. BR data were converted into carbon units using a respiratory quotient of 1.

2.5 Analytical methods

DIN (defined as the sum of nitrate, ammonium and nitrite) was measured by ion chromatography (Metrohm instrument). SRP was determined by spectrophotometry using the ammonium molybdate-potassium antimonyl tartrate method (AFNOR, 2005). DOC concentrations were measured with a total organic carbon analyzer (TOC-L Series, Shimadzu), with a detection limit of 0.01 mg L^{-1} and a precision better than ±5% based on duplicates and standards. Chl*a* concentrations were determined by spectrophotometry after ethanol extraction (90%). Frozen filters were put in 15 mL sterile centrifuge tubes in which 10 mL of ethanol (90%) were added. Tubes were vigorously shaken and then installed in an ultrasonic bath at 70°C for 10 min. Tubes were then stored in a dark chamber for a night, centrifuged 5 min at 4000 rpm and then 10 min at 4000 rpm. Chl*a* concentrations were determined from absorbance at 665 nm after correction of sample turbidity concomitantly measured at 750 nm. Absorbance for CDOM was measured with a Lambda 365 UV/vis spectrophotometer (Perkin Elmer) from 200 to 700 nm (1 nm increment) using a 10 cm quartz cuvette. Napierian absorption coefficients were calculated according to

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$a_\lambda = 2.303 \times A_\lambda / L$,

where a_λ is the absorption coefficient (m^{-1}) at wavelength λ, A_λ the absorbance at wavelength λ and L the path length of the optical cell in m. Spectral slopes for the intervals 275–295 and 350–400 nm were determined from the linear regression of the *log*-transformed a spectra versus wavelength and used to determine the slope ratio (S_R). The slope ratio S_R, calculated as the ratio of $S_{275–295}$ to $S_{350–400}$, is inversely related to the molecular weight distribution of DOM (Helms et al., 2008).

FDOM was measured with a Fluorolog-3 spectrometer (Horiba) using a 1 cm quartz cuvette across excitation wavelengths of 270 – 450 nm (5 nm increment) and emission wavelengths of 300 – 500 nm (2 nm increment) in order to build excitation-emission matrices (EEMs). Lamp intensity and instrument calibration were systematically verified before running samples. EEMs were acquired in sample emission to lamp reference mode, and a correction matrix provided by the manufacturer in both excitation and emission dimensions was automatically applied during acquisition. EEMs were then decomposed into individual components using PARAFAC algorithms (Stedmon et al., 2003). Additional samples collected in Lake Geneva and in the Rhône Basin upstream of Lake Geneva were included in the model (total numbers of EEMs > 800). EEMs preprocessing (Raman scattering removal and standardization to Raman units) was performed prior the PARAFAC modeling. Normalization was done using a Milli-Q water sample run the same day as the sample. A eight components PARAFAC model was obtained using the drEEM 0.3.0 Toolbox (Murphy et al., 2013) for MATLAB (MathWorks, Natick, MA, USA). Split-half analysis, random initialization and visualization of residuals EEMs were used to test and validate the model. The positions of maximums peaks of the PARAFAC components were compared to previous studies with the open fluorescence database OpenFluor using the OpenFluor add-on for the open-source chromatography software OpenChrom (Murphy et al., 2014). The maximum fluorescence F_{Max} values of each component for a particular sample provided by the model were summed to calculate the total fluorescence signal F_{Tot} of the sample in Raman units. The relative abundance of any particular PARAFAC component X was then calculated as $\%C_X = F_{Max}(X)/F_{Tot}$. Precision of EEMs-PARAFAC, based on replicate measurements (n = 5) of different samples, was ±0.001 R.U. for F_{Max} values of components C1 to C5 and ±0.003 R.U for F_{Max} values of components C6 to C8, and ±0.2% for $\%C_X$ for components C1 to C5 and ±0.5% for components C6 to C8. Finally, the variations of PARAFAC components during incubations were expressed as $(F_{Max}(X)_{tf} - F_{Ma}(X)_{t0})/F_{Max}(X)_{t0}$ with t0 and tf the initial and final values of F_{Max}, respectively. Based on the accuracy of EEM-PARAFAC measurements estimated by replicate measurements (see above), we considered variations to be significant if the median of response ratio was higher than ±0.05 for components C1 to C5 and ±0.1 for components C6 to C8.

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253 *2.6 Statistical analyses*

254 Differences in water quality, DOM degradability, BR and consumption of low molecular
255 weight compounds between agro-urban and forest-grassland streams were tested with a Mann
256 Whitney unpaired *t*-test at 0.05 confidence interval level. Differences between the sampling
257 periods were investigated by paired *t*-tests. The level of significance was set to 0.05. The
258 ROUT method implemented in the GraphPad Prism 8 software was used to identify potential
259 outliers. A principal component analysis (PCA) was performed to investigate the importance
260 of human land uses on water quality relative to other geomorphological features (e.g. mean
261 elevation) of the study sites. The data selected for the PCA were DOC, SRP, DIN, and the
262 relative abundance of PARAFAC components collected during the two main campaigns. The
263 PCA was done using the *prcomp* function in the R software, and the *factoextra* package was
264 used to identify the variables that contribute the most to the first two dimensions of the PCA.
265 The sum of cropland and urban areas extents were used as descriptive variables in the PCA
266 biplot.

267 **3. Results**

268 *3.1 Water quality and DOM composition across catchments*

269 Periods of sampling were characterized by relatively similar hydro-climatic conditions
270 with low precipitation and water discharge, and no significant difference was found in the
271 overall water quality between the two campaigns (not shown). However, significant differences
272 were observed between sampling sites depending on their dominant land cover (Figures 2 and
273 3). Chl*a* (total range of variation between sites and periods from 0.2 to 54.9 µg L^{-1}), SRP (from
274 0.7 to 29.3 µg L^{-1}), DIN (from 0.3 to 8.8 mg L^{-1}), and DOC (from 0.7 to 5.9 mg L^{-1}) concentrations
275 were higher in agro-urban streams that were also characterized by DOM of lower molecular
276 weight (higher S_R values) and higher BR values (Figure 2).

277 Eight PARAFAC components were identified in our study sites (Table 2, Supplementary
278 Figure S1), all of which having been already described in previous studies (Graeber et al.,
279 2012; Lambert et al., 2017; Massicotte and Frenette, 2011; Stedmon et al., 2011; Stedmon
280 and Markager, 2005; Williams et al., 2016; Yamashita et al., 2010). Components included three
281 humic-like fluorophores (C1, C3, C4), one fulvic-like fluorophore (C2), two microbial protein-
282 like fluorophores (C5 and C6), and the common tryptophane (C7) and tyrosine (C8) protein-
283 like fluorophores. All components exhibited higher F_{Max} values in agro-urban streams, although
284 the most notable increases were observed for protein-like fluorophores (Figure 3).

285 The two first component of the PCA explained 72.4% of the variance (Figure 4). The
286 first principal component (PC1) was related to protein-like components C6 to C8, DOC and
287 SRP (positive scores) and to humic-like components C2 to C4 (negative scores), whereas the
288 second principal component (PC2) was related to higher C5, DOC and DIN concentrations

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(positive scores) and lower C1 (negative score). The results of the PCA showed that the variability in DOM composition and nutrient loadings was largely driven by land uses (Figure 4). Scores along PC1 were positively related to croplands (Pearson r = 0.49, p = 0.0265), urban areas (Pearson r = 0.63, p = 0.0027) and negatively to forest (Pearson r = -0.72, p = 0.0003) and grasslands (Pearson r = -0.48, p = 0.0295), but not to catchment area (Pearson r = -0.04, p = 0.86), Strahler order (Pearson r = 0.13, p = 0.57) or mean elevation (Pearson r = -0.32, p = 0.18). No relationship was found between PC2 and geomorphological properties of sampling sites, suggesting an in-stream origin for the components C1 and C5. Based on the optical properties of PARAFAC components (Table 2), PC1 represented a shift in the dominant composition of DOM from terrestrial (components C2-C4) to autochthonous (C6-C8) signatures as human disturbance (croplands + urban areas) increases. PC2, however, reflected the in-stream generation of DOM through photodegradation (C1) and bacterial processing of DOM (C5).

3.2 Kinetics of bacterial DOM degradation and consumption of low molecular weight compounds

All incubations were successfully modeled by a first order exponential decay model (r^2 = 0.98±0.02), and we estimated the decay constants and the amounts of BDOC, STRC and LTRC pools for all experiments (Figure 5). BDOC ranged from 0.2 to 2.3 mg L^{-1} (mean = 1.0±0.6 mg L^{-1}), corresponding to 9.7 to 57.6% of initial DOC (mean = 33.8±11%). Higher amount of BDOC in agro-urban streams was accompanied by higher decay constants (from 0.01 to 0.26 d^{-1}, mean = 0.12±0.07 d^{-1}) and greater availability of STRC (from 0.01 to 1.3 mg L^{-1}, mean = 0.5±0.4 mg L^{-1}) but no significant difference was observed regarding the amount of LTRC (from 0.1 to 1.0 mg L^{-1}, mean = 0.5±0.3 mg L^{-1}) across stream categories. Regarding the evolution of the different fractions of DOM during incubations, no significant changes in F_{Max} values were observed for humic-like components during incubations (Figure 6). Component C5, however, exhibited a positive variation in most experiments, showing that this component tended to be produced upon bacterial activity. On the contrary, the other protein-like components C6 – C8 were consumed during incubations. Finally, the consumption of low molecular weights compounds was greater in agro-urban streams for AA and CAR molecules, but no difference was observed regarding the degradation of C&AA (Figure 7).

4. Discussion

The spatial variability in water quality and DOM sources and composition observed in streams and rivers of the Lake Geneva basin echoes numerous previous works illustrating the impact of human activities on freshwater ecosystems (Lambert et al., 2017; Shang et al., 2018; Wilson and Xenopoulos, 2009). Croplands and urbanization resulted in the enrichment in

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protein-like, low molecular weight DOM that was found to enhance DOM bioavailability as well as bacterial respiration. However, we found no evidence of an impact of human activities on the degradation of terrestrial DOM, neither that the latter contributed to ecosystem respiration.

4.1 Origin and biodegradability of DOM in agro-urban streams

DOC concentrations in agro-urban streams of the Lake Geneva Basin were higher compared to those measured in forest-grassland streams (Figure 2) and these increases were accompanied by a shift in DOM composition toward more autochthonous signatures (Figure 4). Effects of agriculture and urbanization on freshwater DOM can vary widely depending on the environmental context and catchment properties (Shang et al., 2018; Stanley et al., 2012), and in our study we attributed this pattern to the combination of enhanced autochthonous production and higher transfer of terrestrial material. Enrichment in nutrients and increased light exposure in agriculture and urban streams can promote primary production (Catford et al., 2007; Stanley et al., 2012; Taylor et al., 2004), and greater algal biomass in our study sites was evidenced by higher Chl*a* concentrations (Figure 2) and the subsequent release of protein-like components (C6 – C8) related to algal DOM (Figure 3, Table 2). Although C5 also relates to autochthonous biological production (Stedmon et al., 2011), its accumulation during incubation experiments implies that this component reflected recently DOM produced by bacterial activity (Figure 6). Higher F_{Max} values of C5 in human-impacted catchments thus represented a positive feedback loop where greater primary production enhanced bacterial activity that shaped DOM composition toward a more bacterial signature (see also Williams et al. 2010). Although of less amplitude, higher F_{Max} values of components associated with terrestrial inputs (C2 – C4, Table 2) and/or photobleaching (C1) indicated a more efficient export of terrestrial material in agro-urban streams. As urbanization tends to limit the hydrological connection between terrestrial and aquatic ecosystems (e.g. Hosen et al. 2014), it is likely that this pattern reflected greater erosion of agricultural soils (Celik, 2005; Graeber et al., 2012). While it was not the purpose of this study to investigate deeper the links between human land uses and DOM sources, our results point that agriculture and urbanization can act in concert to disturb both terrestrial and aquatic sources of DOM leading to greater amount of both humic- and protein-like components in agro-urban streams compared to more natural land cover.

Along with changes in DOM sources and composition, the bacterial consumption of DOM was strongly impacted by land use (Figure 5). Similarly to previous studies carried out in urban catchments (Hosen et al., 2014; Parr et al., 2015), the amount and decay constants of bioavailable DOM increased in human-influenced streams and rivers. However, both autochthonous and terrestrial DOM pools could have contributed to this pattern in the Lake Geneva Basin. The specific reactivity of fluorophores measured during incubations and the

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deconvolution of BDOC into short- and long-term reactive pools allowed us to attribute this increase in bioavailability solely to the consumption of organic compounds generated in stream. Algae are a major source of low molecular weight compounds in aquatic ecosystems through exudation and cell lysis (Kaplan and Bott, 1989), and the release of such labile molecules was evidenced by higher consumption of amino acids and carbohydrates in agro-urban streams (Figure 7) and by the efficient degradation of PARAFAC components related to algal DOM (Figure 6). The strong relationships between the amount of BDOC, the decay constants *k*, and the size of the STRC pool with the initial contribution of protein-like components C6-C8 (Figures 2 and 5, Supplementary Figure S2) provide another evidence that greater DOM bioavailability in agro-urban streams resulted from greater in-stream production.

To the inverse, we found no evidence that the bacterial degradation of terrestrial DOM was influenced by human land uses despites higher contents in inorganic nutrient, higher bacterial activity, and freshly produced autochthonous DOM in agro-urban streams. Terrestrial (C2 – C4) and photo-produced (C1) components showed no significant trends during incubations (Figure 6), yet substantial amount of DOC was consumed during the second phase of the incubation as indicated by the similar sizes of the LTRC and STRC pools (Figure 5). LTRC was positively related to DOC concentrations and F_{Max} values of C1 – C4 components but not with protein-like components (data not shown), implying that this C was somehow related to terrestrial inputs. Although the composition of the molecules that formed the LTRC pools and the drivers of its loss remain unclear, our results evidenced that human land uses did not impact the bacterial consumption of this terrestrial material.

4.2 Linking bacterial respiration to DOM origin and implication of human activities on the transformation of terrestrial DOM along fluvial networks

The loss of terrestrial DOM through bacterial mineralization as it moves along fluvial networks contributes to CO_2 emissions toward the atmosphere (Lapierre et al., 2013) and human activities are expected to impact the role of inland waters in the global carbon cycle by disturbing DOM sources and composition (Xenopoulos et al., 2021). Although limited to the Lake Geneva Basin, our data do not support this assumption. Higher BR in agro-urban streams was indeed mostly related to the accumulation and mineralization of molecules generated in-stream by aquatic primary producers (Figure 8), although the photodegradation of terrestrial DOM could also have contributed to increase BR through the transformation of complex and aromatic molecules into compounds of lower molecular weight (Bertilsson and Tranvik, 1998) as suggested by the positive relationship between BR and C1. The fact that BR relies mainly on DOM derived from primary production in agro-urban streams implies that most of the C released toward the atmosphere as CO_2 by bacterial respiration corresponded to atmospheric CO_2 previously fixed by aquatic producers and converted into biomass. Considering moreover

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that algal DOM can fuel BR also downstream, as suggested by the relationships between C6 – C8 with BR and STRC (defined as the amount of C consumed after 6 days), our results imply that human activities had only a minor effect on the C budget and exchanges between inland waters and atmosphere in Alpine fluvial networks.

Additionally, the consumption of terrestrial DOM did not differ across land use. Moreover, the lack of relationship between LTRC and BR falls in line with experimental studies showing that, in presence of algal derived DOM, terrestrial DOM is preferentially incorporated into bacterial biomass while autochthonous DOM is allocated to respiration (Guillemette et al., 2016). On average, the consumption of terrestrial DOM (i.e., the LTRC pool) at the timescale of our incubation experiments represented 19±8% of initial DOC, an estimation closed to those reported in a medium-size river basin (400 km^2) draining temperate wetlands where only 15% of initial DOC was removed along the whole network (Wollheim et al., 2015). Although we cannot exclude that this terrestrial material could be degraded over longer timescale once entered in the lake Geneva (whose the water residence time is about 12 years), our results are consistent with other studies suggesting that inland waters have only a limited ability to degrade terrestrial C delivered to coastal margins (Borges et al., 2015; Coble et al., 2019; del Giorgio and Pace, 2008; Hanley et al., 2013).

Finally, despite the large range of size of our sampling sites (Table 1), we found no relationship between Strahler order (ranging from 2 to 7) and DOM composition and reactivity. This observation contrasts with a recent study where stream order (ranging from 1 to 4) correlated negatively with humic-like DOM but positively with protein-like DOM (Shang et al., 2018), a pattern consistent with a general conceptual trend describing DOM transformations along fluvial continuum. Indeed, the control of DOM dynamic along the river continuum is expected to shift from a dominant influence of terrestrial inputs in the headwaters to a dominant influence of in-stream removal and autochthonous production as stream order increases (Creed et al. 2015). In our study, however, human land uses had a major role in controlling DOM sources and reactivity at the basin scale (Figure 4). The only exception was a positive correlation between Strahler order and the relative proportion of C5 (Supplementary Figure S3), indicating that the degradation of autochthonous DOM in agro-urban streams led to a bacterial imprint on the DOM pool that persists along the aquatic continuum.

5. Conclusion

In the Lake Geneva Basin, human land uses were found to alter the terrestrial and aquatic sources of freshwater DOM. Greater autochthonous production of DOM in agro-urban streams led to higher amount of bioavailable DOM, stimulating ecosystem respiration while no influence on the loss of terrestrial DOM was observed. Despite their dominant influence on

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DOM composition and reactivity at the basin scale, we found that human land uses had a limited effect in terms of net C flux exchanges between inland waters and atmosphere related to DOM mineralization by heterotrophic bacterial communities.

Enrichment in protein-like DOM due to greater autochthonous production is a recurrent observation in agricultural and urban catchments, therefore the results of this study are likely not limited to the Lake Geneva Basin. However, caution should be taken when extrapolating the impact of human activities on the role of inland waters on the C cycle. Indeed, the net effects of agriculture and urbanization on freshwater DOM vary widely depending on the environmental context (Stanley et al., 2012), leading to apparent opposite effect on BDOC. Thus, while our results are in line with previous works (Hosen et al., 2014; Parr et al., 2015), it contrasts with studies reporting no influence of human land uses on the bacterial consumption of DOM despite higher contribution of autochthonous DOM (Kadjeski et al., 2020; Lu et al., 2013) or higher DOM degradability in agriculture streams related to larger proportion of soil-derived DOM (Shang et al., 2018). Therefore, additional works on the links between human activities and DOM reactivity and fate are needed in order to fully assess the future of inland waters in the context of the global C cycle.

Author Contributions: T. L. conceived the study with contribution from M.-E. P. T. L., P. P., and N. E. collected field samples. T. L. made laboratory analysis. T.L. drafted the manuscript which was substantially commented upon and amended by M.-E. P., P. P., and N. E. All co-authors approved the manuscript.

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669 **Table caption**

670 **Table 1** – Selected properties and dominant classification of sampling sites

671 **Table 2** – Spectral properties (positions of maximum excitation (ex) and emission (em) peaks)
672 of the eight PARARAC components identified in this study, general description and dominant
673 sources based on previous studies.

674

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675 **Figure caption**

676 **Figure 1** – Map of the Lake Geneva Basin and the ten independent catchments sampled
677 during this study.

678 **Figure 2** – Boxplots of (A) Chl*a*, (B) SRP, (C) DIN, (D) DOC concentrations and (E) S_R values
679 and (F) BR in agro-urban (grey) and forest-grassland (white) streams. The box represents the
680 first and third quartile, the horizontal line corresponds to the median, the cross corresponds to
681 the average, and the error bars correspond to the maximum and minimum. Mann Whitney
682 unpaired *t*-test were used to test for statistical differences: ns represents not significant, ∗
683 =p<0.05, ∗∗ =p<0.01, ∗∗∗ =p<0.001, ∗∗∗∗ =p<0.0001.

684 **Figure 3** – Boxplots of F_{Max} values of PARAFAC components in agro-urban (grey) and forest-
685 grassland (white) streams. The box represents the first and third quartile, the horizontal line
686 corresponds to the median, the cross corresponds to the average, and the error bars
687 correspond to the maximum and minimum. Mann Whitney unpaired *t*-test were used to test for
688 statistical differences: ns represents not significant, ∗ =p<0.05, ∗∗ =p<0.01, ∗∗∗ =p<0.001, ∗∗∗∗
689 =p<0.0001.

690 **Figure 4** – PCA biplot, including loadings plot for the input variables and scores plot for
691 stations. Markers are shaped according to the sampling period and colored according to a
692 gradient of human disturbance (defined as the sum of % croplands and % urban areas,
693 supplementary ordinal variable in the PCA).

694 **Figure 5** – Boxplots of (A) BDOC concentrations, (B) constant decay *k*, (C) STRC and (D)
695 LTRC pools in agro-urban (grey) and forest-grassland (white) streams. The box represents the
696 first and third quartile, the horizontal line corresponds to the median, the cross corresponds to
697 the average, and the error bars correspond to the maximum and minimum. Mann Whitney
698 unpaired *t*-test were used to test for statistical differences: ns represents not significant, ∗
699 =p<0.05, ∗∗ =p<0.01, ∗∗∗ =p<0.001, ∗∗∗∗ =p<0.0001.

700 **Figure 6** – Response ratio of PARAFAC components during incubation experiments with tf =
701 day 28 and t0 = day 0. Grey bars represent threshold of significance above or below which
702 significant production or consumption of component was identified. See text for details.

703 **Figure 7** – Boxplots of (A) AA , (B) CAR, and (C) C&AA consumption in agro-urban (grey) and
704 forest-grassland (white) streams. The box represents the first and third quartile, the horizontal
705 line corresponds to the median, the cross corresponds to the average, and the error bars
706 correspond to the maximum and minimum. Mann Whitney unpaired *t*-test were used to test for
707 statistical differences: ns represents not significant, ∗ =p<0.05, ∗∗ =p<0.01, ∗∗∗ =p<0.001, ∗∗∗∗
708 =p<0.0001.

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709 **Figure 8** – Relationships between BR and (A) DOC concentrations, (B) the sum of initial F_{Max}
710 values of C6, C7, and C8 components, and (C) initial F_{Max} values of C1 component

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712 Table 1

713

Site	Area (km^2)	Mean elevation (masl)	Strahler order	Forest (%)	Croplands (%)	Urban areas (%)	Classification
1 - La Combe	39	1241	2	73	3	1	Forest-grassland
2 - Le Boiron	11	692	2	62	36	1	Forest-grassland
3 - Le Grand Curbit	14	614	2	24	70	4	Agro-urban
4 - La Venoge	228	696	4	31	60	6	Agro-urban
5 - La Mèbre	21	597	3	18	38	43	Agro-urban
6 - La Sorge	12	595	3	24	65	11	Agro-urban
7 - La Dranse	638		4	33	14	5	Forest-grassland
8 - Le Rhône	5238	2124	7	23	2	3	Forest-grassland
9 - La Veveyse	65	1105	5	45	20	6	Forest-grassland
10 - La Paudèze	16	775	4	33	40	25	Agro-urban

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715 Table 2

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Component	Max ex (nm)	Max em (nm)	Description & dominant sources
C1	<270	424	Widespread humic-like fluorophore, terrestrial[a,b] and/or photoproduced[c].
C2	<270 (330, 380)	498	Fulvic-like fluorophore, widepread, terrestrial origin[d,e].
C3	<270 (355)	438	Humic-like fluorophore, widespread, terrestrial origin[d].
C4	320	402	Low molecular weight humic-like fluorophore, related to agriculture[a,b,e].
C5	300	336	Protein-like fluorophore associated with biological production[f].
C6	<270	372	Anthropogenic humic-like fluorophore related to algal[g] or bacterial[f] production in urban areas[e,g].
C7	275	332	Tryptophan-like fluorophore, indicative of autochthonous production[b,c].
C8	270	304	Tyrosine-like fluorophore, indicative of autochthonous production[b,c].

[a] Stedmon & Markager, 2005; [b] Yamashita et al., 2010; [c] Massicote & Frenette 2011; [d] Graeber et al., 2012; [e] Lambert et al., 2017; [f] Stedmon et al., 2011; [g] Williams et al., 2016.

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718 Figure 1

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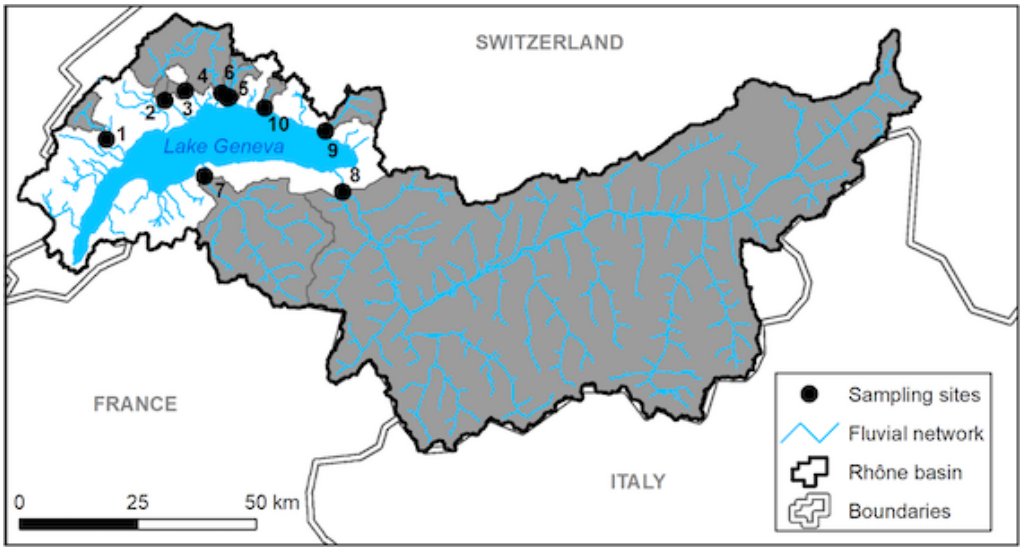
720



721

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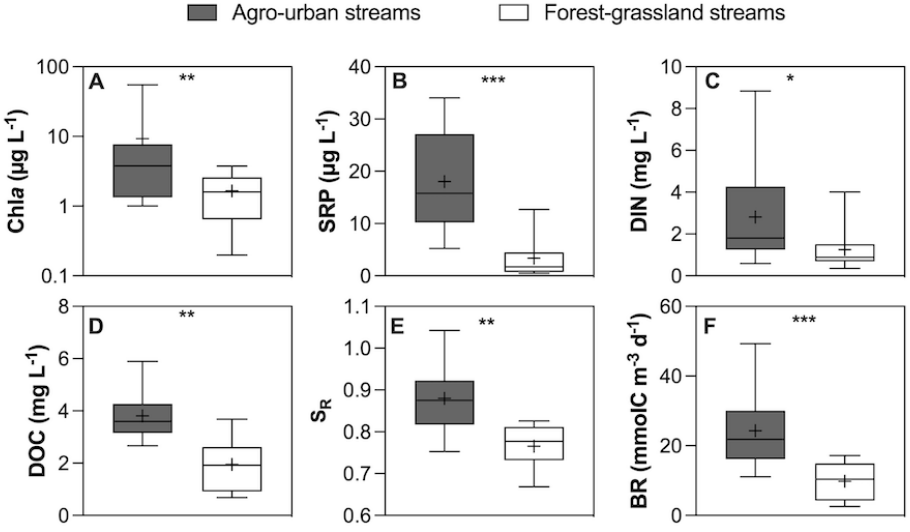
722 Figure 2



723

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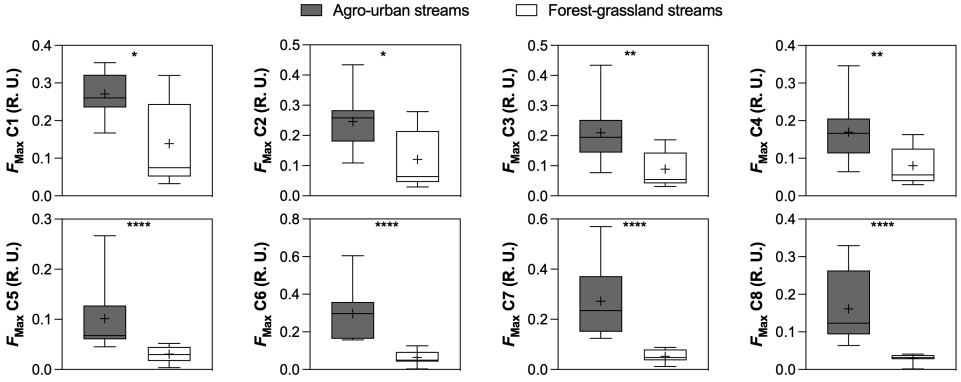
724 Figure 3



725

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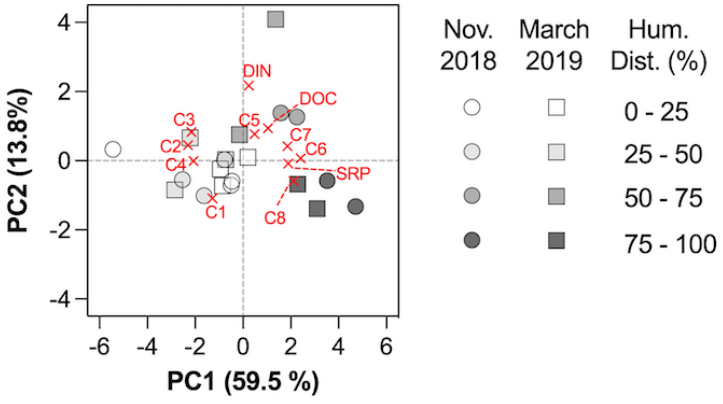
726 Figure 4



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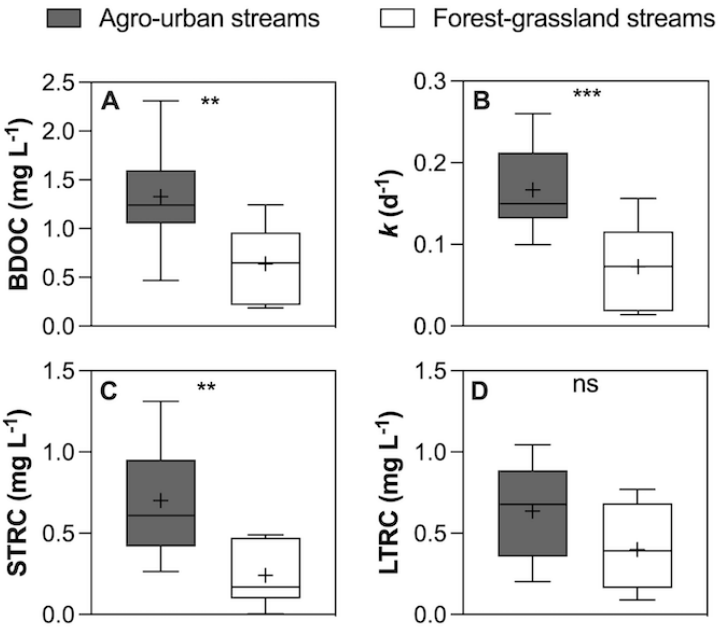
728 Figure 5



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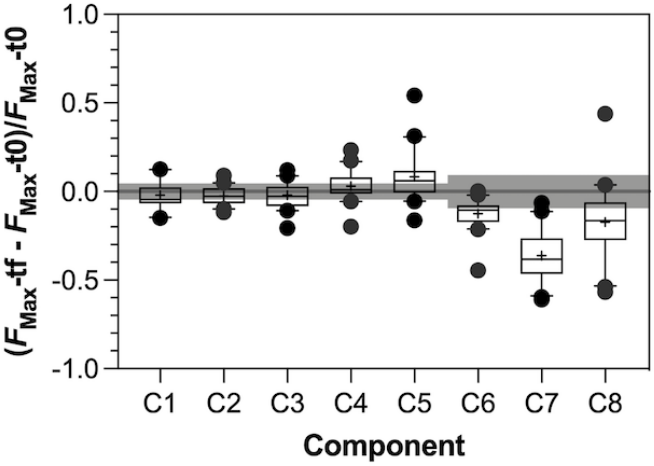
730 Figure 6



731

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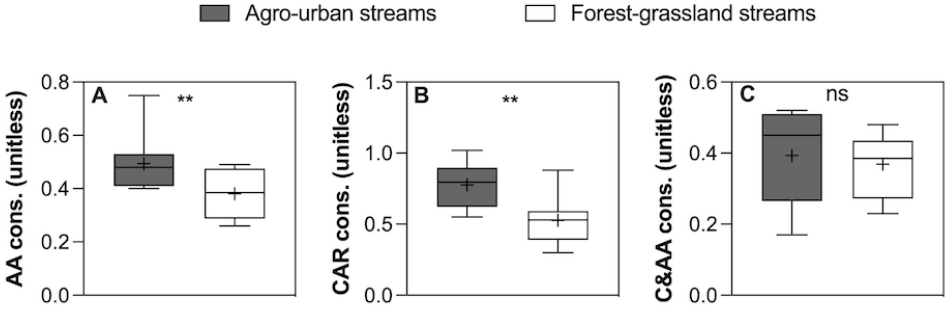
732 Figure 7



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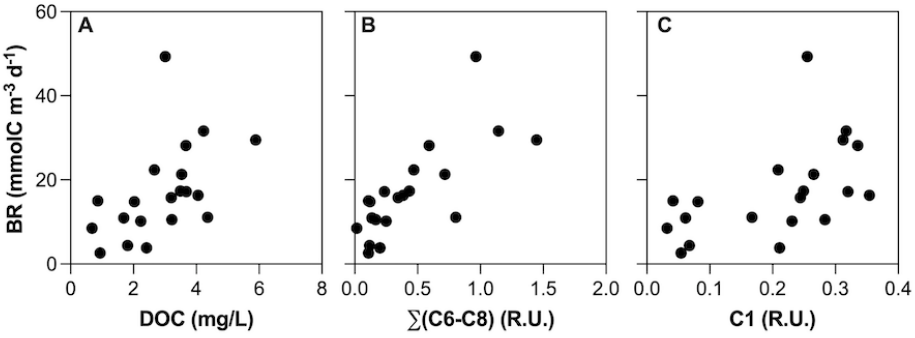
734 Figure 8



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