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Seawater copper content controls biofilm bioaccumulation and microbial community on microplastics

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Abstract. The understanding of the interactions between trace metals and microplastics (MPs) is hampered by the poorly understood role of biofilm in trace metal bioaccumulation. In this study, the influence of seawater copper content on biofilm microbial communities' composition as well as on the resulting Cu biofilm bioaccumulation on MPs was investigated within seawater of Toulon-bay (NW Mediterranean Sea), highly impacted by anthropogenic pollution. Based on incubation experiments, the bioaccumulation of copper (Cu) was examined following an increasing gradient of its concentrations. Two sets of incubation experiments were run with seawater supplied with MPs, sampled in two sites in which Cu contents differed: Pt12 (most contaminated site) and Pt41P (less contaminated site). For each incubation experiment, 5 treatments were considered differing in Cu concentrations, ranging between 30 and 400 nM and between 6 and 60 nM, for Pt12 and Pt41p, respectively. A control experiment (filtered at 0.2 μm) was run in parallel for each incubation experiment. At the time scale of the incubation period, both prokaryotic and eukaryotic richness and diversity of the formed biofilm onto MPs was higher in the more contaminated site, thus appearing as the most driving factor. In addition, Cu addition clearly shaped biofilm communities, evidencing co-occurrence patterns between prokaryotes and eukaryotes with diatoms playing a central role. Remarkably, these differences in biofilm formation were reflected in the amount of bioaccumulated Cu per dry weight of MPs, thus exhibiting higher values in the most contaminated site. Within this site, the increase of Cu seawater content enhanced its bioaccumulation onto MPs until reaching saturation. This study strongly suggests a striking link between seawater copper content, biofilm community shaping and the resulting Cu bioaccumulation onto MPs.

Keywords: Microplastics; bioconcentration; copper; microbial community composition; biofilm.

1. Introduction

Over the last few decades, plastic debris of the millimetre size range (<5 mm), identified as microplastics (MPs) (Derraik, 2002; Andrady, 2011, Law, 2017), have become of an increasing concern due to their prevalence within the water column (114 studies performed in the last 20 years; Aretoulaki et al., 2020), impact on ecosystems (Huang et al., 2020; Khalid et al., 2020) and their potential to sorb chemical contaminants (Holmes *et al.*, 2012; Turner and Holmes, 2015; Brennecke *et al.*, 2016; Acosta-Coley *et al.*, 2019). Hydrophobic organic pollutants have been highlighted as being incorporated into MPs and/or adsorbed onto their surface (Fossi *et al.*, 2012; Antunes *et al.*, 2013). In contrast, sorption of trace metals (TMs) on MPs as well as their incorporation into these artificial particles have been shown as limited (Klößner et al., 2020; Zou et al., 2020).

Nevertheless, these reported studies on TMs sorption/incorporation were performed on pristine MPs, therefore on polymer matrices devoid of any biological colonization. However, MPs are clearly suitable for biofilm formation as they were shown to host a diverse microbial community, including heterotrophs, autotrophs, predators and symbionts, referred to as 'a plastisphere' (Zettler *et al.*, 2013; Amaral-Zettler et al., 2020). Because biofilms are typically built up by a complex matrix of extracellular polymeric compounds, it is very likely that its formation on MPs could increase and

modify the exchange surface of these synthetic particles within the water column, which would potentially influence the interaction with TMs. Therefore, TMs load on MPs should be looked at whilst considering the role of biofilm in this accumulation. So far, to the best of our knowledge, only few studies involving MPs, biofilm and metals were conducted, thus revealing a higher metal sorption when comparing MPs as pristine *versus* colonized by biofilm (Guan et al., 2020). Recently, metal accumulation has been shown to correlate with the amount of biofilm (Richard et al., 2019). Although evidence emerged supporting the role of biofilm in enhancing the adsorption and stabilization of trace metal on MPs (Wang et al., 2020), the magnitude of their bioconcentration on MPs as well as the feedback influence on its microbial community composition is still poorly known.

Toulon bay is a coastal marine ecosystem in the NW Mediterranean Sea (South of France), representative of an area coping with anthropogenic metallic pollutants mainly attributed to an important anthropization and extensive harbours activities. Concentrations of TMs in Toulon-bay have been reported as being of the same order of magnitude or even higher than strongly impacted aquatic ecosystems (Pringault *et al.*, 2012; Misson et al., 2016), with a decreasing gradient from the most anthropized parts to the open waters of the Mediterranean Sea. Along this gradient, bacterioplankton composition is shaped, including shifting the autotrophic community from picoeukaryote dominated community in the most contaminated parts to cyanobacteria (more representative of the open ocean) in less impacted areas (Coclet et al., 2018, 2019, 2020). Alongside TMs pollution, an important amount of MPs can be found in the surface water of Toulon bay, with an average of 32 800 MPs/km⁻² (Tesan-Onrubia et al; 2021). Potential bioaccumulation of TMs on MPs may also influence microbial biofilm diversity. Recently, biofilm formation on plastics panels did not appear to be mainly driven by seawater TMs content (Coclet et al., 2021, Catao et al., 2021a), despite strong evidence of the influence of Cu on

other biofilm communities (Catao et al., 2019, 2021b). However, in the Toulon-bay as well as in other coastal areas, a better understanding of the interactions of MPs-TMs is critical as MPs are most likely to be ingested by marine organisms and potentially transferred to high trophic levels through the trophic network (Andray, 2011).

In this study, incubation experiments with natural seawater were conducted to investigate the relationship between an increasing range of seawater Cu content and biofilm formation MPs, its composition including both prokaryotes and eukaryotes, and its role in Cu bioaccumulation.

2. Methods

2.1. Seawater sampling and bioassay experiments

In this study, two sets of incubation experiments were carried out with surface seawater sampled into 10-L LDPE carboys at two locations in the bay of Toulon, Pt12 (43.11 °N; 5.93°E) and Pt41p (43.26°N; 6.58°E) (Fig. 1). Sampled seawater was then filtered *in situ*, through 90 µm mesh plankton net and homogeneously distributed once in the laboratory, into 1 L ultraclean PFA bottles, under a laminar flow hood. Experimental sampling and incubation bottles were previously washed three times with Milli-Q water (18.2 MΩ, Millipore), then with 10% (v/v) nitric acid (HNO₃, Analytical Grade, Fisher), rinsed three times with Milli-Q water then filled with Milli-Q water acidified with 0.1% HNO₃ Trace Metal Grade (Fluka). For all experiments, PFA bottles were chosen so as to prevent any metallic contamination.

The sampling locations were chosen based on the long-time monitoring of trace metals performed in the area, highlighting contrasted Cu concentrations in the considered locations (Tessier et al., 2011, Layglon et al., 2020; Coclet et al., 2018, 2019, 2020). Pt12 sampling station is located in the enclosed part of the bay of Toulon, near the main harbour,

therefore submitted to important metallic inputs from the anthropogenic activities (Fig. 1). In contrast, Pt41p sampling station is located in the large bay of Toulon, with trace metal concentrations similar to the encountered values in the Mediterranean Sea. In both sampling stations, subsamples were taken for *in situ* measurements of Cu and exhibited concentration of 30 nM for Pt12 and 6 nM and Pt41p.

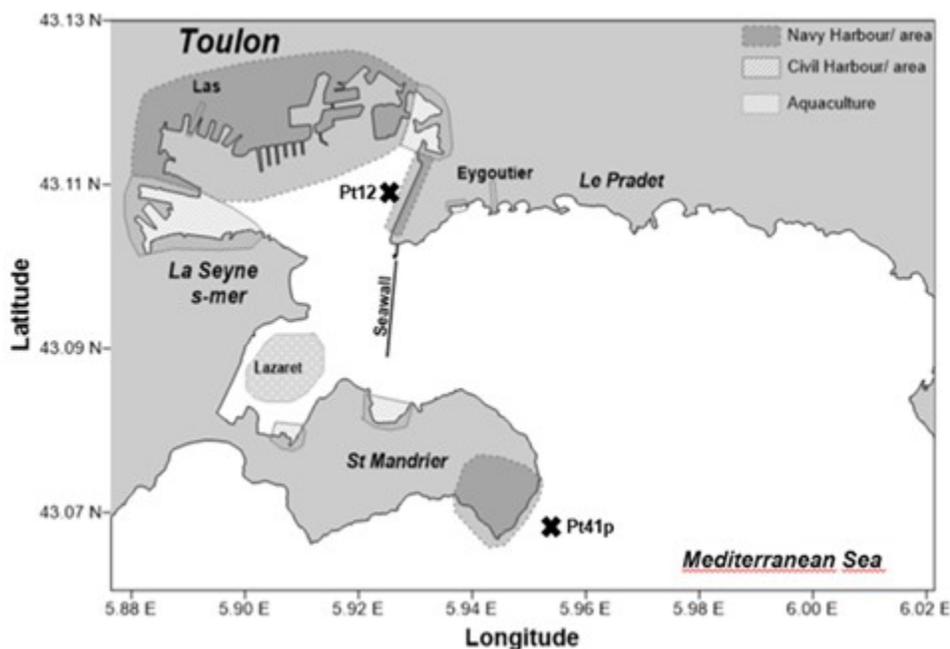


Figure 1: Sampling stations in the small (Pt12) and large (Pt41p) bay of Toulon.

The MPs pellets used in this study were obtained by flash freezing of polymeric polyethylene microbeads (CRT102,50, Carat) in liquid nitrogen and their subsequent grinding, using an ultra-centrifugal mill (ZM200, Retsch). Obtained grinded pellets were further sieved from 500 to 100 μm and were thoroughly rinsed with Milli-Q water. MPs were then dried (1 h, 60°C) and pellets within a size range between 200 and 300 μm were further used.

The two sets of experiments were run in triplicates (1-L each). Seawater from Pt12 or Pt41p was incubated with a single density of MPs (10 mg L⁻¹) and an increasing gradient of Cu concentrations. Two ranges of Cu concentrations were applied whether the seawater was sampled in Pt12 or Pt41p as the initial natural Cu concentrations differed: 30 and 6 nM in Pt12 and Pt41p, respectively. Thereby, Cu was added to reach final concentrations of 60, 100, 200 and 400 nM in the seawater sampled at Pt12 (hereafter, small bay experiment) and of 20, 30 and 60 nM in the seawater sampled at Pt41p (hereafter, large bay experiment). For each sampling station, a triplicate without any Cu addition (reflecting in situ concentration), was run in parallel. Likewise, two bottles of 1-L of sterile (filtered at 0.2 µm, precleaned acetate filters) seawater amended with 10 mg L⁻¹ MPs and a Cu concentration of 200 nM and 60 nM for seawater of Pt12 and Pt41p, respectively, were also run as controls. All experimental bottles were gently mixed, using PFA stirring bars and were incubated over 13 days at 18°C, in a phytotron (Mlr 351 Sanyo) with a day night cycle of 12:12 h.

2.2. Subsampling over the incubation period

Incubated seawater was subsampled at T₀ (1 h after MPs and Cu amendment), an intermediate time (T₇ or 8), and at T_{final} (after 13 days of incubation period) for bacterioplankton cell abundance and only at T_{final} for Cu. Subsamples for Cu (10 mL) were filtered through 0.2 µm precleaned acetate syringe filters (HCl, trace metal grade), acidified with 0.5% HCl (trace metal grade) and stored at 4°C until analysis. Subsamples for heterotrophic and autotrophic cell abundance determination were fixed with 0.5% glutaraldehyde (final concentration), incubated in the dark for 15 min at ambient temperature and then transferred into -80°C freezer, until analysis. Unfiltered seawater (10 mL) was subsampled at the same time scale for both heterotrophic and autotrophic bacterioplankton cell abundance. These samples were fixed with 0.5% glutaraldehyde (final

concentration), incubated in the dark for 15 min at ambient temperature and then transferred into -80°C freezer, until analysis.

At the end of the incubation period, for all experiments, each incubation bottle was filtered through 0.2 µm polycarbonate filters (HCl washed, trace metal grade) to recover MPs. Half of the recovered MPs was used for biofilm diversity determination, through DNA extraction while the other half was suspended in 10 mL NaOH (0.1 M) and mixed up for 24 h, for biofilm extraction. A few randomly chosen pellets were also fixed for biofilm visualization using confocal laser scanning microscopy (CLSM).

2.3. DNA extraction, amplification, sequencing, and data processing

DNA from MPs was extracted using the DNeasy PowerBiofilm Kit (Qiagen, Courtaboeuf, France) according to the manufacturer's instructions. For all samples, the V4-V5 region of the 16S rDNA gene was targeted and amplified using 515F-Y/926R primers (Parada et al., 2016) for prokaryotes identification whereas the 18S rDNA gene was targeted and amplified using V7 primers (Debroas et al., 2017) for eukaryotes identification. Amplicons were sent to Eurofins (Konstanz, Germany) for Illumina MiSeq 2 × 300 pb paired-end sequencing.

Sequences were analysed following the FROGS process (Escudié et al., 2018) in a Galaxy environment following current guidelines. Clustering relies on Swarm which does not agglomerate sequences based on the typical 97% threshold but relies instead on both the number of differences and the likely series of accumulation of those differences (Mahé et al., 2014). Chimera detection relies on VSEARCH with *de novo* UCHIME method (Edgar et al., 2011; Rognes et al., 2016). The innovative cross-sample validation step from FROGS was used to confirm the chimeric status on all samples. Rare OTUs representing less than 0.005% of all sequences were removed. Affiliation by blastn+ was performed using the SILVA v132

database for prokaryotes and the Protist Ribosomal 2 (PR2) database (Guillou et al., 2012) for eukaryotes. The datasets for this study can be found in NCBI SRA database, Bioproject PRJNA735079.

2.4. Confocal laser scanning microscopy (CLSM) and Scanning Electron microscopy (SEM)

MPs were washed in artificial seawater and subsequently fixed with 3.8% formaldehyde solution. To visualize cells, staining was performed with 250µl of Syto 9 or Syto 61 (1µM-ThermoFisher) in 24-well plates for 30 min, in the dark and at room temperature. In addition, exopolymeric substances (EPS) characterizing biofilm development were also visualized using FITC (proteins), concanavalin A (ConA, sugar) and calcofluor white (CFW, sugar). After 2 washes in phosphate buffered saline solution, microplastics were mounted on microscope slides with one drop of Prolong Diamond antifade (ThermoFisher) and incubated for 24 hours at room temperature. Confocal images were performed with 20X/0.75NA and 63xoil/1.4NA objectives, on a Zeiss Confocal LSM 510 Meta (excitation: 488 nm - emission: BP 505-530 nm).

2.5. Copper extraction and analysis

NaOH mixtures were filtered (0.2 µm) and the retained MPs were dried and then weighed. A volume of 5 mL of the filtered NaOH mixture was taken and completed to 10 mL with filtered Milli-Q water (18.2 MΩ, Millipore). Then, the solution (10 mL) was UV-irradiated (Metrohm) for 24 h in presence of H₂O₂, to ensure a complete digestion of the extracted biofilm. The solution was finally analysed for Cu concentration.

Cu analyses were performed by cell-inductively coupled plasma mass spectrometry (ICP-MS; Perkin Elmer NexIon 300X), in a direct injection mode. The international geostandards SLRS-3 (Riverine water reference material) was used to check the validity and reproducibility of the analysis.

To avoid interference with seawater salt, incubation bottle samples were diluted 25 times in nitric acid, while extracted biofilm samples were diluted only 5 times. With obtained Cu concentrations in the incubation bottle (T=13 days) and in the extracted biofilm, the partition coefficient (Kd) was determined as follow:

$$Kd (L/Kg) = ([Cu] \text{ Biofilm} / \text{MPs mass}) / [Cu] \text{ final}$$

Where Cu biofilm and Cu final are Cu concentrations in the extracted biofilm solution and in incubated water after 13 days of incubation time, respectively. MPs mass is the dry weight of MPs.

2.6. Flow cytometry analyses

Cell abundance counting was performed using a FACSCalibur (BD Biosciences) flow cytometer (PRECYM flow cytometry platform, <http://precym.mio.univ-amu.fr/>). Instrument specific beads were used to calibrate the cytometer. For heterotrophic bacteria, frozen samples were thawed at room temperature and subsamples were stained with 1:10 diluted SYBR Green II (2 μ L, Molecular Probes) and incubated for 15 min in the dark, in order to stain nucleic acids. Each cell was then characterized by 3 optical parameters: light diffusion side scatter (SSC); green fluorescence (515-545 nm; SYBERgreen), and red fluorescence (670 LP; chlorophyll-a). Group-specific phytoplankton were distinguished based on plots of forward scatter vs. orange fluorescence (phycoerythrin containing *Synechococcus* like), and forward scatter vs. red fluorescence (eukaryotes). Size classes of eukaryotic phytoplankton were further distinguished based on forward scatter (pico, nano and large eukaryotes). Flow cytometry data were acquired and counted with the Cell Quest Pro software.

2.7. Statistical analysis

Cu concentrations in the incubation bottles as well as in the extracted biofilm, Kd and alpha diversity indexes were compared using a one-way analysis of variance (ANOVA) and Tuckey posthoc tests (p-value > 5%).

Alpha diversity indexes (Chao 1, Shannon), clustering using Weighted Unifrac distances (Prokaryotes) and Unweighted Unifrac distances (Eukaryotes) followed by Principal Coordinates Analysis (PCoA) and PERMANOVA were performed using R Cran version (3.5.0.4) with “phyloseq”, “vegan” and “ggplot” program packages. The approach taking only into account the presence or absence of eukaryotic taxa, in addition to phylogenetic distances, was considered appropriate to account for possible biases associated with the limited quantification capacity of eukaryotic metabarcoding (Kelly et al. 2014). SIMPER analyses were performed to precise the contributors to the dissimilarity between pairs of samples (PRIMER® 6).

Interactions among prokaryotic and eukaryotic OTUs were identified using CoNeT (Co-occurrence Network) plugin 1.1.1.beta (Faust et al., 2012). Data were processed using Pearson and Spearman correlation, with Bray-Curtis and Kullback-Leiber dissimilarity distances and using a threshold of 150 and 400 edges for Pt12 and Pt41p, respectively. Edge specific p-values were computed according to Brown’s method, in the permutation step. In the final step, the Benjamini-Hochberg multiple testing correction was applied ($p < 0.05$).

3. Results

3.1. Influence of Cu seawater content on pico- and nanoplankton cell abundance within the incubation experiments

One hour after Cu amendments (T0), picophytoplankton composition in seawater from Pt12 (Fig. S1-a) was dominated by picoeukaryotes (between 6000 ± 427 and 11000 ± 364 cells mL⁻¹), followed by

Prochlorococcus-like (between 508 ± 12 and 1630 ± 158 cells mL⁻¹) and *Synechococcus*-like (between 45 ± 4 and 603 ± 40), in all considered treatments. Following the increasing gradient of Cu (from 30 to 400 nM), the initial cell abundance of each group significantly decreased (ANOVA; $p < 0.01$). After 13 days of incubation (for each experiment, bars on the right, Fig. S1-a), *Synechococcus*-like disappeared while both picoeukaryotes and *Prochlorococcus*-like drastically decreased (ANOVA; $p < 0.01$), particularly in treatments with the highest levels of Cu. Initial nanophytoplankton cell abundance (for each experiment, bars on the left, Fig. S1-c) in Pt12 exhibited a decrease with the increase in Cu gradient. After 13 days of incubation, nanophytoplankton cells abundance significantly increased (ANOVA; $p < 0.01$) compared to T0, in treatments having received 30 and 60 nM of Cu, remained constant with amendments of 100 and 200 nM of Cu (ANOVA; $p > 0.05$) and decreased in the treatment with 400 nM of Cu (ANOVA; $p < 0.01$). Regarding heterotrophic bacteria (Fig. S1-e), initial cell abundance was not influenced by the increasing Cu gradient. On the contrary, after 13 days of incubation time, cell abundance significantly decreased in each treatment (ANOVA; $p < 0.01$). Nevertheless, following the increase in Cu concentration, an increase in final cell abundance (T=13 days) was observed, except in the treatment with 400 nM (ANOVA; $p < 0.01$) (Fig. S1).

In seawater from Pt41p (large bay of Toulon), initial picophytoplankton abundances were higher than at Pt12 (Fig. S1-b) and the composition here was dominated by *Synechococcus*-like, followed by *Prochlorococcus*-like and finally picoeukaryotes (ANOVA; $p < 0.01$), in each treatment. As for seawater from Pt12, after 13 days of incubation period, cell abundance dropped in all treatments (ANOVA; $p < 0.01$), within each group. This decrease was more pronounced for treatments with the highest Cu concentrations. The initial nanoplankton cell abundance was, conversely to picophytoplankton, lower at Pt41p than at Pt12 (Fig. S1-b). Initial cell

abundance significantly decreased in treatments with 30 and 60 nM of Cu (ANOVA; $p < 0.01$). After 13 days of incubation time, the nanoplankton abundance trend was not clear. Regarding heterotrophic bacteria (Fig. S1-f), the initial cell abundance remained similar (ANOVA; $p = 0.38$) among treatments. After the incubation period, for treatments with Cu concentration lower than 20 nM, the bacterial cell abundance slightly increased (ANOVA; $p < 0.05$) compared to T0, while it decreased for Cu concentrations of 30 and 60 nM (ANOVA; $p < 0.01$).

In both investigated sites, principal component analysis (PCA) on cell abundance of the different identified groups showed that picoplankton composition in mesocosms was more influenced by the incubation experiment period rather than Cu seawater content (Fig. S2).

3.2. Biofilms visualization on MPs

The visualization of microbial cells using CLSM (Fig. 2, Fig. S3) showed that microbial colonization settled on MPs in incubation experiments conducted at both Pt12 and Pt41p whereas MPs incubated with 0.2 μm sterile filtered seawater (control treatments) did not show any microbial colonization. In addition, using specific markers, exopolysaccharide (EPS) were also detected within the biofilm formed on MPs, thus showing enriched regions in proteins (FITC) or sugar (concanavalin A and calcofluor white) (Fig. 2).

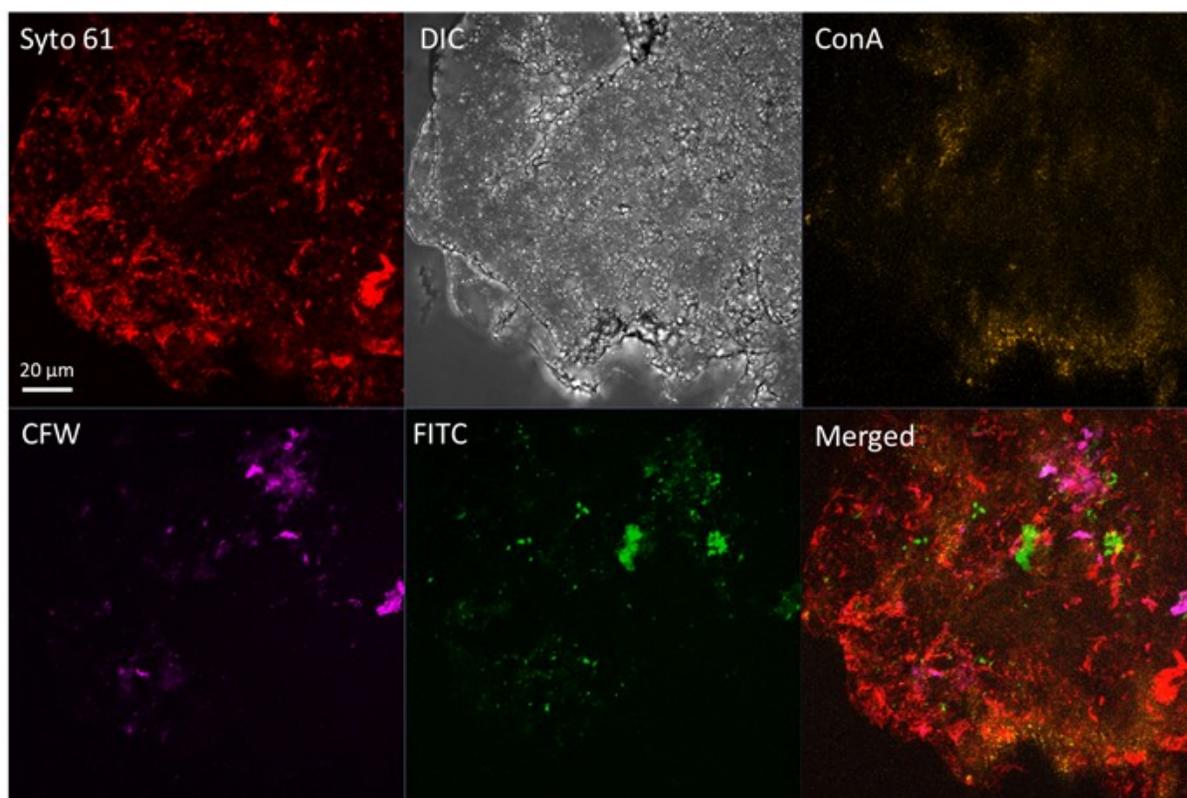


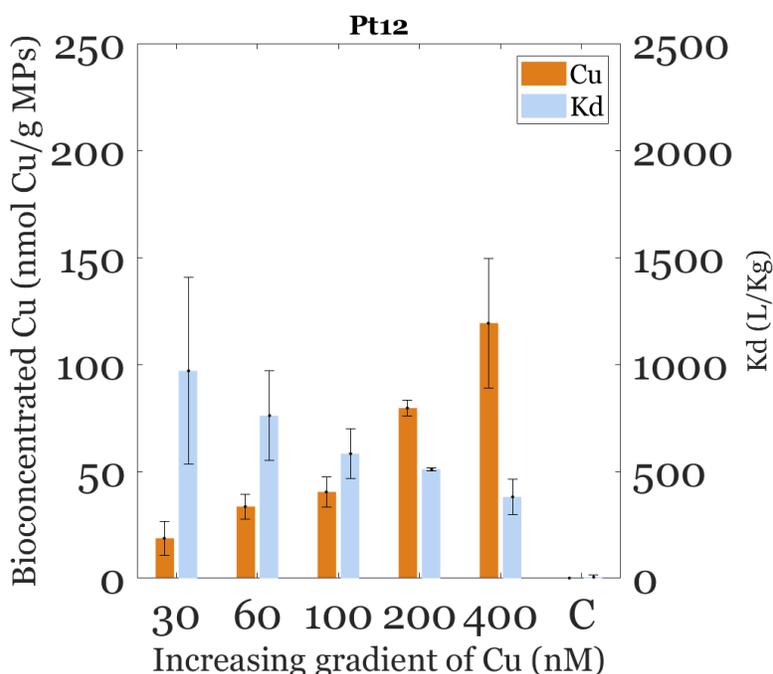
Figure 2: Confocal Laser Scanning Microscopy of microplastics. Attached microbial cells were visualized using the SYTO 61 as well as differentiated interference contrast microscopy (DIC). In addition, exopolymeric substances (EPS) characterizing biofilm development were visualized using FITC (proteins), concanavalin A (ConA, sugar) and calcofluor white (CFW, sugar).

3.3. Effect of an increasing gradient of Cu on biofilm and its bioaccumulation on MPs

In the water from Pt12 (small bay of Toulon), following an increasing gradient of Cu concentrations (between 30 and 400 nM, Fig. S4), the amount of bioaccumulated Cu per dry weight of MPs exhibited a significant increase (*ANOVA*; $F=23.3$; $df=4$; $p<0.01$), with values ranging between 18 ± 7 and 119 ± 30 nmol/g MPs (Fig. 3). Conversely, although not significantly different due to the variability within triplicates (*ANOVA*;

$F=3.1$, $df=4$, $p=0.07$), K_d seemed to exhibit a decrease of its average value, from 970 ± 437 to 380 ± 83 L kg⁻¹ for treatments with 30 and 400 nM of Cu, respectively. In the sterile filtered treatment, both bioaccumulated Cu (ANOVA; $F=31.9$; $df=4$; $p<0.01$) and K_d (ANOVA; $F=7.7$; $df=4$; $p<0.01$) were low (Fig. 3).

In the water from Pt41p (large bay of Toulon), the amount of bioaccumulated Cu per dry weight MPs was similar between treatments (ANOVA; $F=0.8$; $df=4$; $p=0.53$), ranging between 0.52 ± 0.06 nmol/g MPs and 1.06 ± 0.06 nmol/g MPs (Fig. 3). Similarly to the previous case, K_d exhibited an decrease with the increasing gradient of Cu (ANOVA; $F=5.9$; $df=4$; $p=0.02$), ranging between 12 ± 10 Kg L⁻¹ and 66 ± 15 Kg L⁻¹.



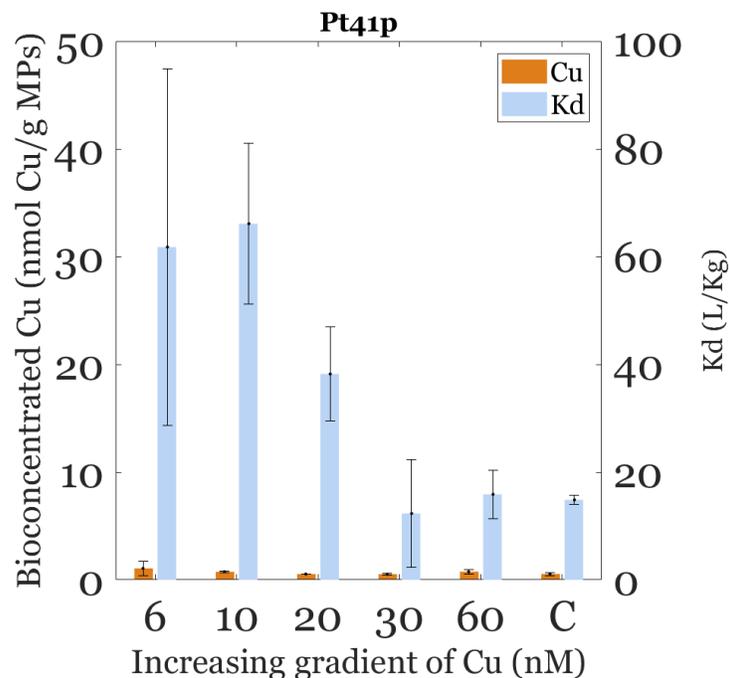


Figure 3. Bioaccumulated Cu on microplastics at the scale of the incubation period and its associated repartition coefficient (Kd) at Pt12 (small bay) and at Pt41p (large bay). C refers to treatments with sterile filtered seawater. Note that the scale of y axes of both bioaccumulated Cu and Kd are different at Pt12 compared to Pt41p.

3.3. Alpha diversity of microbial biofilm on MPs

The Illumina Miseq run produced 1984472 and 2340978 paired-end reads, leading to 1543558 and 2018175 merged sequences for 16SrDNA and 18SrDNA gene sequencing respectively. 1030 and 512 OTUs could be generated after clustering for prokaryotes and eukaryotes respectively. Archaea were excluded from the prokaryotic set of reads (<0.5%). Some samples could not be amplified due to low DNA extracts (Pt41p_C2r1 and r3, C3r1 and r2, C4r3 for 16S rDNA and Pt41p_C5r3 for 18S rDNA). Sequence normalization by a random resampling to the sample with the fewest sequences was performed (n=13345 and 24308 for prokaryotes and

eukaryotes respectively), leading to the exclusion of few samples (Pt12_C5r3 for 16S rDNA and Pt41p_C1r2 and C3r2 for 18S rDNA).

All alpha diversity indexes (Fig. 4) displayed significant differences among biofilms developed on MPs, within seawater from the two sites (Pt12 and Pt41p) and for both prokaryotes and eukaryotes. Richness and diversity were always much higher at Pt12 (small bay) than at Pt41p (large bay). Considering the influence of Cu seawater content on prokaryotes, only the highest concentration (400 nM) decreased the Chao 1 at Pt12 and both Chao 1 and Shannon index at Pt41p, when compared to natural seawater. No significant difference could be noticed for eukaryotes regardless of the index.

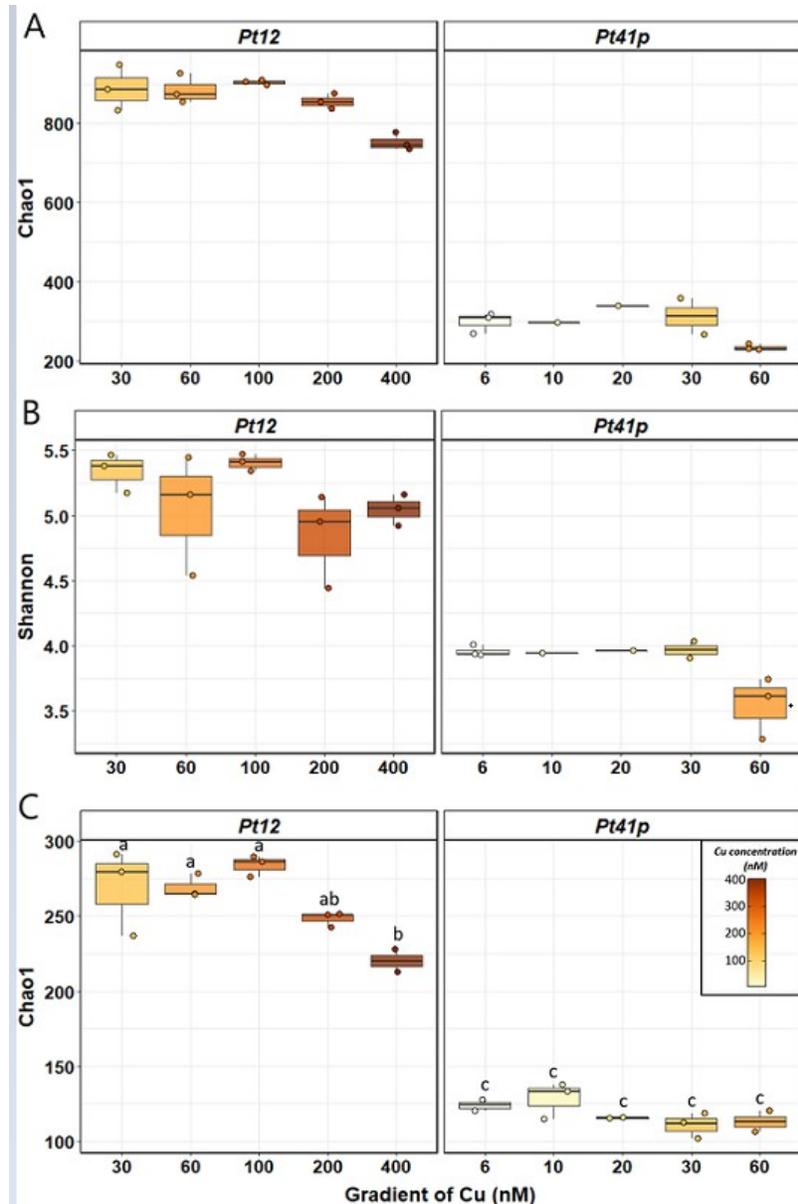


Figure 4: Alpha diversity indexes for experiments at both sites Pt12 and Pt41p: Chao1 index (A) and Shannon index (B) for prokaryotes and Chao1 index for eukaryotes (C).

3.4. Beta-diversity and composition of microbial biofilms on MPs

For both prokaryotic and eukaryotic communities, the first clustering factor was definitely the site (Fig. 5). In addition, in each incubation experiment, a clear shift of natural communities was observed on the y-axis following the

increasing gradient of Cu, except for eukaryotes from Pt41p for which no dissimilarity could be noticed (Fig.5; panel B).

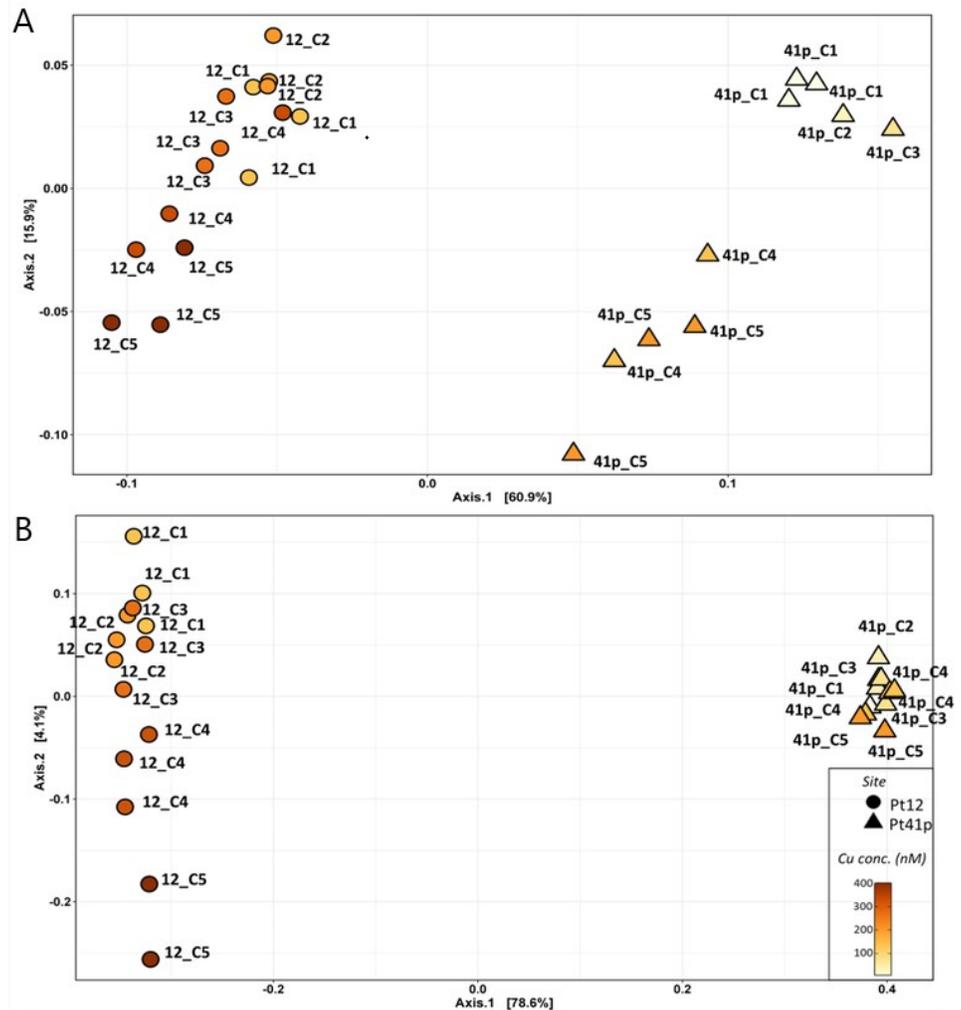


Figure 5: Principal Coordinates Analysis (PCoA) of biofilm prokaryotes (A) and eukaryotes (B) from microplastics, for incubation experiments conducted in the seawater from the Pt12 (circles) and Pt41p (triangles). Clustering was studied using Weighted Unifrac distances for prokaryotes and Unweighted Unifrac distances for eukaryotes.

Prokaryotic biofilm communities were dominated at the two sites by three classes: alpha-, gammaproteobacteria and Bacteroidia. The dissimilarity between communities from the two sites was shown at the family level with communities dominated by Cryomorphaceae (mainly

Luteibaculum, 5 to 30%) and Rhodobacteraceae (mainly *Tateyamaria*, 22 to 32%) at Pt12, but Flavobacteriaceae (mainly *Aureicoccus* and *Pseudofulvibacter*, 28 to 43%) and Sphingomonadaceae (mainly *Erythrobacter*, 7 to 14%) at Pt41p (Fig. 6A and S5). SIMPER analyses showed that the prokaryotic taxa the most involved in the dissimilarity between mesocosms with natural seawater at Pt12 (C1=30 nM) and with 200 nM of Cu (C4) were two OTUs affiliated to *Luteibaculum* (Cryomorphaceae), an unknown Rhodobacteraceae and *Croceibacterium* (Flavobacteriaceae), with the three first being enriched in Cu (Table S1). Taxa the most involved in the dissimilarity between mesocosms filled with natural seawater at Pt41p (D1) and with 60 nM of copper (D4) were four OTUs affiliated to Flavobacteriaceae. *Aureicoccus* and *Winogradskyella* showed higher relative abundance in D1 whereas *Pseudofulvibacter* and *Croceibacter* in D4 (Table S1). In addition, an unknown Rhodobacteraceae and two Sphingobacteriaceae, *Spongibacter* and *Erythrobacter*, highly contributed to the dissimilarity between D1 and D4 and were more abundant in D4 too.

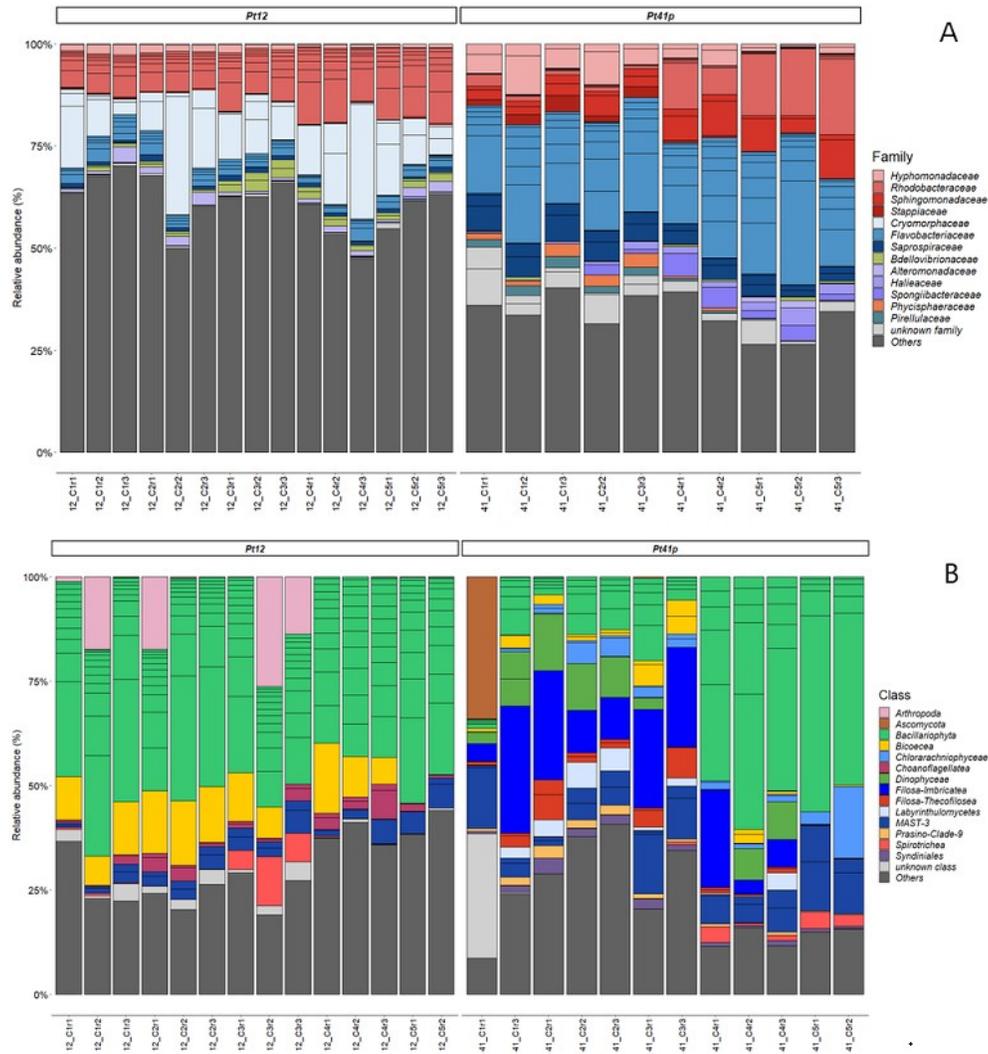


Figure 6: MPs biofilm community composition in the experiments with the seawater from the two sites (Pt12 and Pt41p) at the family level for prokaryotes (A) and the Class level for eukaryotes (B).

Biofilm eukaryotic communities were mostly composed of microeukaryotes (Fig. 6B and S6). *Oithona*, a planktonic copepod (crustaceans), ubiquitous and abundant in oceans, which was probably entrapped in biofilms when filtered, was the only multicellular organism identified, and only in Pt12 samples without copper concentration exceeding 100 nM. Microeukaryotes were dominated by diversified Bacillariophyta at Pt12 (41 to 78%), whereas only samples with the two

highest copper concentrations (30 and 60 nM) exhibited similar dominance at Pt41p. Raphid pennates strongly dominated at Pt12 irrespective of the copper concentration (25 to 57%). Among them, *Cylindrotheca closterium* or *Pseudo-Nitzschia* appeared to be sensitive to copper whereas *Nitzschia* or *Surirella* were enriched in high copper concentration (from 4 to 23% and 1 to 6%, respectively). Similarly, araphid pennates, without taxonomic affiliation at the genus level, were enriched with Cu addition, both at Pt12 (from 2 to 13%) and Pt41p (6 to 16%, even if the percentage decreased just for the highest copper concentration). Polar centric Mediophyceae exhibited the same trend at Pt41p (3 to 44%).

Bicoecea (unicellular flagellates, Stramenopiles) accounted between 8 to 20% at Pt12 and from 2 to 8% at Pt41p, except at the highest Cu concentration for the two sites (less than 1%). MAST-3J (Stramenopiles) represented from 1 to 9% at Pt12, with a decrease at the highest Cu concentration, whereas a higher percentage was observed at Pt41p (7-20%) without any copper effect. Unidentified centroheliozoa completed the community at Pt12, with a clear decrease at the highest Cu concentration (from 8 to 1%). Dinophyceae, including *Pentaparsodinium* that could be present as cysts in biofilms, represent generally around 2% but accounted for up to 13% of the reads at Pt41p with a decrease for the two highest Cu concentrations. Picoplanktonic phototrophs in the Prymnesiophyceae accounted for up to 6% at Pt41p, showing a sensitivity to the highest copper concentration too. Moreover, Filosa Imbricateae and Thecofilosea (Rhizaria) appeared to be also sensitive to copper, with percent decreasing from 30 and 16, respectively, to 0% at the highest copper concentration at Pt41p. Finally, Fungi accounted for less than 2% (except one sample at Pt41p), without clear variation depending on the site or copper concentration, as well as Ciliophora (from 0 to 11%).

SIMPER analyses showed that the eukaryotic taxa was the most involved in the dissimilarity between mesocosms with C1 and C4, where

OTUs mostly affiliated to Bacillariophyta but with different taxa abundances in the two conditions (Table S1, 7). C4 was used rather than C5 because the latter appeared less representative of natural ecosystems. Two OTUs of *Nitzschia*, one *Amphiprora* and an additional but unaffiliated Raphid-pennate contributed to the dissimilarity between conditions and were abundant in C4 whereas *Pseudo-Nitzschia* and an additional but unaffiliated Raphid pennate were more abundant in C1. The copepod *Oithona*, the flagellate *Bicosoeca*, the ciliate *Holosticha* and a Choanoflagellata were the other contributors to the dissimilarity between C1 and C4. The most involved taxa in the dissimilarity between D1 and D4, and enriched in D4, were mostly OTUs affiliated to Bacillariophyceae (Polar centric Mediophyceae, araphid and raphid pennates), whereas taxa abundant in D1 were heterotroph microeukaryotes (*Filosa imbricatea*, Sordariomycetes, centroheliozoa or MAST-3J).

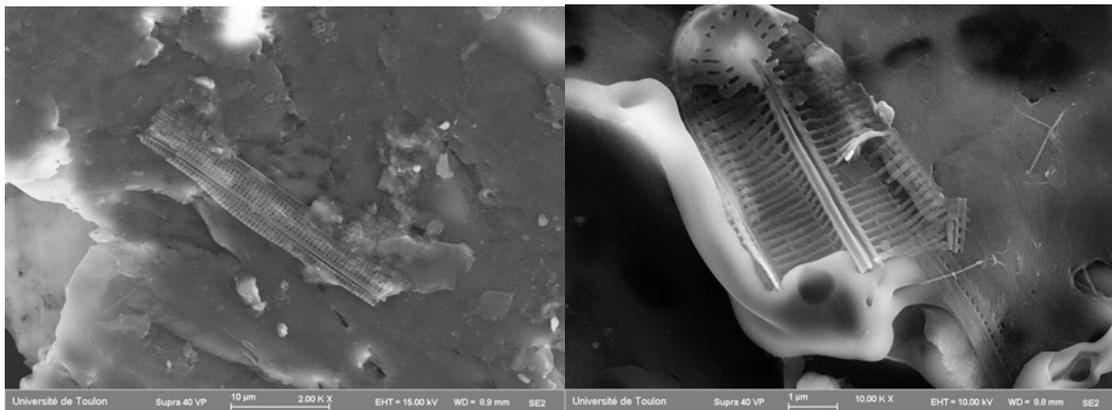


Figure 7: Visualization of diatoms within the biofilm developed onto MPs at Pt12 incubation experiment, using scanning electron microscopy.

Networks allowing to identify the co-occurrence among prokaryotic and eukaryotic taxa on MPs were constructed using CoNet for each of the two experiments (Fig. 8, Table S2). For Pt12, the main cluster of the network displayed 79 nodes (OTUs) associated with 95 degrees or edges. Ten small clusters with only two to three nodes completed the network. Most of the edges were positive (62%). The ten nodes with the highest

number of edges corresponded to five Rhodobacteraceae (*Candidatus Rhodobacter lobularis*, *Pelagimonas*, *Roseobacter*, *Tateyamaria* and an unknown genus) followed by five Bacillariophyta, four of them being araphid pennates including *Grammatophora undulata*. For Pt41p, the main cluster of the network displayed 45 nodes (OTUs) associated with 79 edges. Twelve small clusters with only two to four nodes completed the network. Most of the edges were positive (72%). The ten nodes with the highest number of edges corresponded to three bacterial and seven eukaryotic OTUs showing the highest diversity. Bacteria belonged to the Flavobacteriaceae (*Winogradskyella*), Stappiaceae (unknown genus) and Pirellulaceae (*Blastopirellula*) when Eukaryotes were Bacillariophyta (*Thalassiosira profunda* and an unknown polar centric Mediophyceae) but also three Cercozoa (one unknown genus among Filosa Thecofilosa and another one among Filosa Imbricata, and a mixotrophic amoeboid-like microalgae *Minorisa*), a Prymnesiophyceae (*Phaeocystis*) and a Ciliophora (unknown genus among Strombidae).

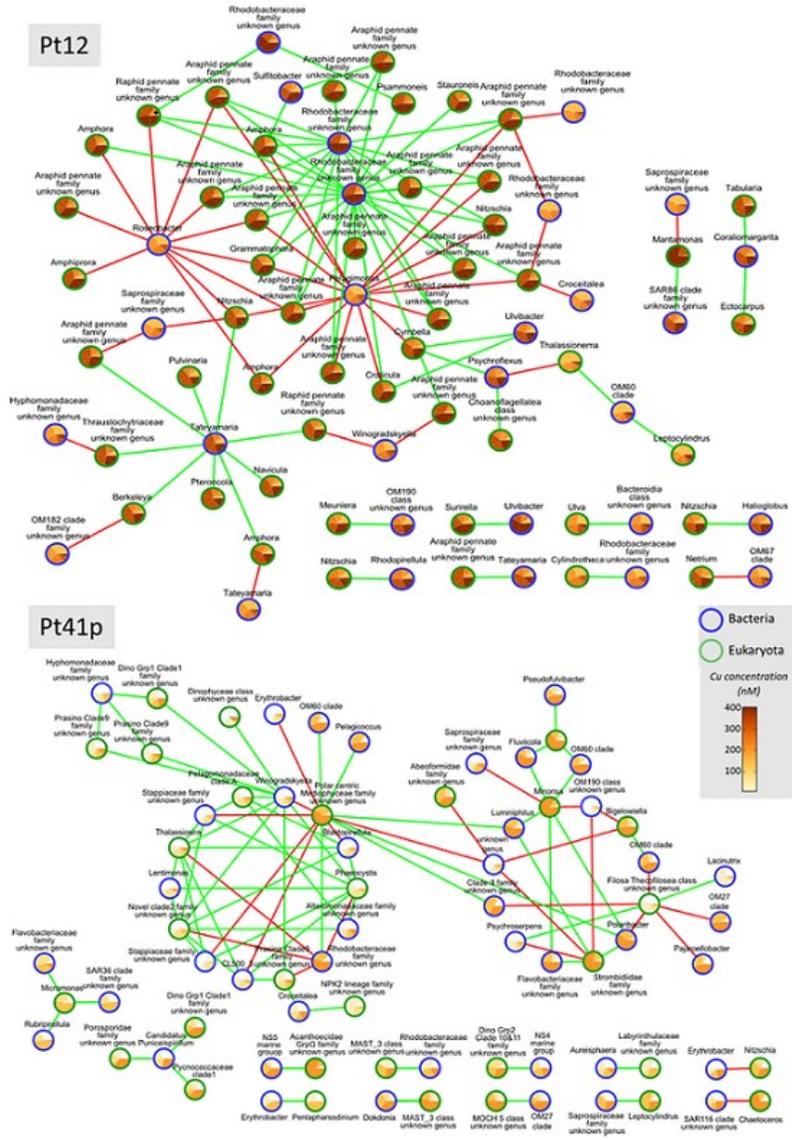


Figure 8: Interactions among prokaryotic and eukaryotic OTUs were identified using CoNET (Co-occurrence Network) in the two experiments with seawater from the sites Pt12 and Pt41p. Blue circles stand for bacterial OTUs whereas green OTUs stand for eukaryotes. Relative abundances in each sample are indicated within circles. Samples are colored depending on the Cu content. Positive edges are in green whereas negative ones are in red.

4. Discussion

In this study, for the sake of viability of encountered microbial communities, Cu was added in a range between 30 and 400 nM and between 6 and 60 nM when seawater was sampled from Pt12 (small bay of Toulon) and from Pt41p (large bay of Toulon), respectively, as Cu concentrations were shown to exhibit a decreasing gradient from the most enclosed part of the bay to the open waters (Coclet et al., 2018, 2021; Layglon et al., 2020). In agreement with *in-situ* observations (Coclet et al., 2018, 2019; Delpy et al., 2018), distinct picophytoplankton composition was observed in each incubation experiment, thus exhibiting picoeukaryotes as a dominant phytoplankton group in seawater from Pt12 (small bay of Toulon) and cyanobacteria (*Synechococcus* and *Prochlorococcus* groups) in seawater from Pt41p (large bay of Toulon). At the time scale of the incubation experiment, all phytoplankton groups as well as heterotrophic bacteria decreased overall following the increasing gradient of Cu, thus supporting the role of Cu in shaping planktonic microbial community composition. Considering this context, this study aimed to demonstrate, using controlled conditions with natural seawater, that: (1) Cu concentrations in seawater controlled its bioaccumulation onto MPs, with this bioaccumulation strictly linked to biofilm formation and (2) Cu bioaccumulation shaped the microbial community within the biofilm formed on MPs.

4.1. Copper bioaccumulation on MPs is due to biofilm

In this work, abiotic alterations of MPs in link with light exposure and mixing (see method section) may have promoted adsorption of Cu onto their surface at the time scale of the incubation period. Yet, the low observed values in 0.2 μm sterile filtered seawater treatments in both incubation experiments highly suggested the role of biofilm as a striking link within MPs-Cu interplay. Trace metals (TMs) bioaccumulation onto MPs through

biofilm formation is still poorly resolved, especially in seawater. As a matter of fact, so far, most studies which reported MPs-TMs interaction in marine and/or freshwater systems focused on pristine microplastics, devoid of any biological colonization, but also on the influence of aging, resulting from physical alterations and/or photooxidation, on these pellets in promoting TMs adsorption (Turner and Holmes 2015; Brennecke et al., 2016; Mao et al., 2020). Nevertheless, there is increasing evidence that investigating the interaction between TMs and pristine MPs can be in fact misleading. Recently, reported studies have shown that biofilm developments on these synthetic particles eased metals accumulation, including Cu (Richard et al., 2019; Guan et al., 2020, Wang et al., 2020).

In this study, biofilm formation around MPs was proven in both studied sites. In addition, dissimilar alpha and beta diversity was observed in biofilms depending on the sites, but also Cu seawater content. This differences in biofilm formations between sites were reflected in Cu bioaccumulation, thus exhibiting quantitatively higher values in seawater from Pt12 (up to 119 ± 30 nmol/g MPs) than Pt41p (up to 47 ± 32 nmol/g MP), with a clearer increasing bioaccumulation of Cu following seawater Cu contents in Pt12. Higher alpha-diversity combined to higher DNA extracts (data not shown) suggested that greater biofilm biomass occurred on MPs at Pt12, in agreement with higher Cu bioaccumulation. Biofilm accumulation of metals have been essentially highlighted in freshwater systems (Ancion et al., 2010; Tien and Chen, 2013), with retention time up to 14 days before a subsequent release into the surrounding waters (Ancion et al., 2010). Within the biofilm, the sorptive properties of EPS have been shown to effectively bind large amounts of metals (Decho, 2000 and references therein). In this study, EPS among others, may have promoted Cu bioaccumulation. In fact, in the water, Cu speciation is dominated by organic complexes (Wong et al., 2018 and reference herein), especially in areas where dissolved organic matter (DOM) is found at high

concentrations. TMs cations complexation by organic ligand or organic matter occurred through several complexation acidic sites (Luo et al., 2009), defined by their concentration and complexing constant. Such parameters are closely related to the surrounding DOM. For example, the complexing affinities towards Cu of a natural DOM considered, from a river end-member to seawater end-member, was revealed to change according to the location along an estuary: from a concentrated one-type of complexing site with a modelled concentration and a high constant to a 2-types of complexing sites with lower concentrations but higher constants (Louis et al., 2009). Similarly, organic ligands produced in the biofilm matrix, EPS for example, will differ from one incubation experiment to another, as richness and diversity indices differ among biofilms. The concentrations and complexing constants of each produced biofilm could not be determined, the dedicated experiments needing an important quantity of organic material (Louis et al., 2009), which is impossible when dealing with the very thin biofilm layer around MPs. Yet, it is well known that a biofilm will produce organic ligands whose nature and quantity depends on the biofilm composition (Fang et al., 2014; Mahapatra et al., 2020). It was already demonstrated that the biofilm cultivated on polyethylene molecularly differs from a biofilm developing on cobblestones, with a weaker interaction between EPS and Cu in the polyethylene biofilm (Wang et al., 2020). It is therefore obvious that the biofilm developing in each of the tested conditions (and especially seawater) of this study will have its own composition and hence organic ligands properties. This is reflected in copper accumulation.

4.2. Biofilm community structure is primarily controlled by the sampling site

Sampling site clearly appeared as the primary driver for biofilm community shaping. Even though Cu seawater contents were significantly different at the two sites, Cu alone may not explain differences in biofilm

microbial community. In fact, a recently conducted study at these two sites, as well as at three additional ones in the Toulon bay, has concluded that the Cu gradient failed to explain the clustering of prokaryotic community structure of mature biofilms (Coclet et al., 2021). In this same study, interestingly, most of the reported environmental biofilm community drivers, such as temperature, pH, dissolved oxygen, and nutrients, were measured and did not show any significant difference among sites, suggesting that additional parameters like DOM, hydrodynamics or top-down control, among others, could have played a role in shaping biofilm microbial community (Catao et al., 2019; Mazoyer et al., 2020; Ferguson et al., 2021). In other studies, performed on biofilms formed on large artificial polymer surfaces, environmental parameters also clearly drive biofilm communities which was reported in the literature too (e.g. Oberbeckman et al., 2016; 2018; Lawes et al., 2016; Briand et al., 2017; Catao et al., 2021). When samples within the plastisphere failed to be related to environmental factors characterizing the sampling site (e.g. Bryant et al., 2016, Dussud et al., 2018), the buoyant character of plastic promoting the influence of multiple environmental conditions may provide an explanation.

In addition, while supplying seawater from Pt41p and Pt12 with the same concentration of Cu (60 nM), biofilm communities on MPs did not converge neither for prokaryotes nor eukaryotes, thus confirming that Cu did not primarily drive biofilm microbial communities. However, the well-known toxicity influence of Cu on phytoplankton can indirectly influence biofilm, through an exerted control on cell abundance in one hand but also via alteration of DOM nature in the other hand. Finally, we assume that the two clusters of biofilm communities observed in mesocosms of the two sites were also the result of different initial planktonic communities. As a matter of fact, biofilm taxa come from the rare planktonic community, particularly copiotrophs, which make their determination within the bacterioplankton

difficult, using classical Illumina approach (Zhang et al., 2019, Catao et al., 2021, Coclet et al., 2021).

Communities on MPs included typical biofilm taxa. Discrepancies for taxa affiliation between this study and the previous biofilm study on polymer surface in the bay of Toulon (Briand et al., 2017) should probably be associated with the considered surface size of plastic (large polymers versus MPs), the type of conducted experiments (mesocosm versus field study), the immersion time (13 *versus* 30 days) but perhaps the interannual variability too. Cryomorphaceae taxa enriched in Pt12 biofilms seem more predominant in coastal ecosystems, with enhanced organic loads (Bowman, 2020), and also enriched in the marine plastisphere (Oberbeckmann et al., 2016). The co-occurrence network analysis highlighted the central role of several Rhodobacteraceae in interactions with eukaryotes. In addition to the low richness and diversity observed in biofilms from Pt41p, taxa reputed as pioneer ones, like *Alteromonas* (Dang and Lovell, 2000; Lee et al., 2008; Pinto et al., 2019) or Hyphomonadaceae (Dang and Lovell, 2016), could be identified, which strengthen our hypothesis of weak biofilms on MPs with seawater from Pt41p.

Considering the poorly studied eukaryotes on MPs (Kettner et al. 2019), most of them were microeukaryotes, as multicellular one's development should be limited by the size of MPs and the incubation time. Unfortunately, the taxonomic affiliation of the 18SrDNA gene (Whatever the data bank PR2 or Silva, data not shown) allowed the identification of only a few of OTUs to the genus level in biofilms on MPs (41% for PR2). Diatoms (Bacillariophyta), especially pennates, clearly dominated the most anthropized site (Pt12) showing a high diversity of genera among the raphid pennate (*Amphiprora*, *Cylindrotheca*, *Navicula*, *Nitzschia*, *Pseudo-Nitzschia*, *Surirella*). Likewise, the co-occurrence network underlined the significance of diatoms in interactions with prokaryotes and primarily with Rhodobacteraceae. Despite often identified on scanning electron microscopy (*e.g.* Zettler et al. 2013,

Kirstein et al., 2018), diatom communities were scarcely studied on MPs compared to prokaryotes. Diatoms in biofilms were studied on large artificial surfaces, all using microscopic approaches, mainly in tropical areas, and with the aim to describe their diversity depending on the substrate type including antifouling coatings (Patil and Anil, 2005, Mitbavkar and Anil, 2008, Molino and Wetherbee, 2008, Zargiel et al., 2014, Briand et al., 2017, Muthukrishnan et al., 2017). This major microphytobenthic taxa was conversely highly investigated in freshwater ecosystems, mostly in the context of bioindication (*e.g.* Sabater et al., 2017, Rivera et al., 2018, Polst et al., 2018).

Pt41p exhibited the highest number of families with a lower importance of diatoms in the natural seawater condition. The presence of vagile predators of bacteria like unicellular flagellates Bicoecea or amoeboids-like Filosa Imbricateae and Thecofilosea could be explained. However, diatoms are mainly represented by polar centric mediophyceae including *Chaetoceros*, Dinophyceae including *Pentapharsodinium* or picoplanktonic phototrophs in the Prymnesiophyceae, which are more surprising to be found in biofilms. Specifically considering the centric diatoms, some authors tend to consider them as members of the plastisphere, but considering their preferential planktonic lifestyle and taken in consideration that specific lifestyle occurred for other microorganisms as bacteria (*e.g.* Wright et al., 2020), we may hypothesized that such polar centric diatoms may belong to the rare biofilm community. Consequently, a contamination issue with planktonic taxa from the mesocosms seawater during the filtration of the MPs should not be excluded. Moreover, the high number of polar centric mediophyceae, which increased with Cu addition in biofilms on MPs especially in Pt41p, could be also linked to the relative increase of nanophytoplankton observed in the mesocosm seawater (Figure S1-2).

4.3 Cu also controlled biofilm community on MPs

As mentioned above, Cu was not identified as the main driver of biofilm community on plastic surface in a previous experiment with natural seawater Cu concentrations in Toulon bay (Coclet et al., 2021). Alpha-diversity appeared not affected in this study as well except at the highest Cu concentration (400 nM), which though is not environmentally realistic as such high concentrations are not naturally observed. Conversely, Cu clearly influenced the beta-diversity of biofilm communities for each site, except for eukaryotes with seawater from Pt41p. It may look contradictory because substrates were immersed in waters from the same sites, and conventional plastics (polycarbonate and polyethylene in these two cases) are now reputed to exhibit similar biofilm communities (Kirstein et al., 2018, Wright et al., 2020). However, the size of the substrates was significantly different with 5x5 cm coupons in the previous experiment (Briand et al., 2017) whereas MPs were used here. Hence, thicker biofilms have probably grown on MPs (Fazey and Ryan, 2016), allowing higher interactions between copper and biofilm cells, consequently with a higher selection within the biofilm communities compared to thick and mature biofilms observed on large substrates. The substrate size effect was still not fully studied but CLSM pictures confirmed the presence of thin biofilms on MPs.

Biofilms on MPs formed with the seawater from Pt12 exhibited higher richness, diversity and Cu bioaccumulation than at Pt41p, which suggested that a higher biomass, including EPS. While some EPS were clearly visualized with CLSM, their more in-depth study and characterization may provide a better understanding of interactions with metals. As two dissimilar biofilm communities occurred with seawater from the two sites, the assumption of dissimilar produced EPS can be made. However, the higher resulting Cu bioaccumulation in Pt12 compared to Pt41p could also relate to a quantitatively higher EPS production rather than the production of qualitatively different EPS. Even reputedly complex to study, EPS would be a key component for further insight regarding the interaction between

biofilm and metal bioaccumulation (Flemming et al., 2016). The relative contribution of bacteria and diatoms to this production will be another important key point to address as well.

The Rhodobacteraceae genus *Tateyamaria* which was observed as one of the most enriched taxa at Pt12 following Cu content increase here, was already enriched in the most contaminated site in the previous study in the Toulon bay (Coclet et al., 2021). In addition, the network analysis highlighted the co-occurrence of this genus with several diatoms (*Amphora*, *Berkeleya*, *Navicula*, *Nitzschia*, *Pteroncola*), especially in high Cu concentration. Nevertheless, it remains difficult to conclude on simple co-selection, direct or indirect interactions. *Tateyamaria* was also reported as a biomarker on copper-based antifouling paints, with a specific correlation with the abundance of the copper resistance gene CueO (Catao et al., 2021). Similarly, the relative abundance of *Erythrobacter* (Sphingomonadaceae) increased in biofilms from Pt41p with the increasing copper content, which is consistent with its enrichment in biofilm community on copper-based antifouling coatings (Catao et al., 2019, 2021). As already reported for copper resistance in the context of antifouling paints, the selection was mainly observed at the genus or species level rather than family. As a matter of fact, especially considering two of the main families, Rhodobacteraceae and Flavobacteriaceae, it was already reported that both sensitive and tolerant to copper taxa were identified in the same biofilm communities (Catao et al., 2019, 2021).

Among *Bacillariophyta*, *Cylindrotheca closterium* or *Pseudo-Nitzschia* appeared to be sensitive to Cu whereas *Nitzschia* or *Surirella*, but also *Amphora* or *Grammatophora* to a lesser extent, were enriched in high Cu concentrations. Similarly, araphid pennates, without taxonomic affiliation at the genus level, were enriched with Cu addition, both at Pt12 and Pt41p. The selection of resistant taxa together with the decrease of sensitive ones indicated that, similarly as for freshwaters, community structure and/or

specific taxa could be used for bioindication (Tlili et al., 2016, Vasselon et al., 2017). Indeed, Bacillariophyta could be used to find out if MPs move through contaminated or non-contaminated sea waters, which could give information on MPs -followed current.

5. Conclusion

This study revealed a tight interaction among MPs, biofilm and Cu concentration in seawater of the bay of Toulon (NW Mediterranean Sea). Indeed, Cu bioaccumulation on MPs was clearly linked to biofilm. Furthermore, the seawater Cu-content controlled biofilm bioaccumulation on MPs but was also involved in biofilm microbial communities shaping.

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