How should ecohazard of micropollutants in wastewater be gauged? Using bioassays to profile alternative tertiary treatments

Matteo Papa1,2,*, Lidia Paredes1*, Donatella Feretti3,6, Gaia Viola3, Giovanna Mazzoleni4,6, Nathalie Steinberg4,6, Roberta Pedrazzani5,6, Juan Lema1, Francisco Omil1, Marta Carballa1

1Department of Chemical Engineering, Institute of Technology, Universidade de Santiago de Compostela, Rua Lope Gomez de Marzoa, E-15782 Santiago de Compostela, Spain
2Department of Civil, Environmental, Architectural Engineering and Mathematics, University of Brescia, via Branze 43, I-25123 Brescia, Italy
3Department of Medical and Surgical Specialties, Radiological Sciences and Public Health, University of Brescia, Viale Europa 11, I-25123 Brescia, Italy
4Department of Clinical & Experimental Sciences, University of Brescia, viale Europa 11, I-25123 Brescia, Italy
5Department of Mechanical and Industrial Engineering, University of Brescia, via Branze 38, I-25123 Brescia, Italy
6Brescia University Research Center “Integrated Models for Prevention and Protection in Environmental and Occupational Health” (MISTRAL), Italy

*These authors contributed equally to this work.

Abstract

The research on emerging pollutants in wastewater has become a worldwide issue of increasing environmental concern, especially considering the growing interest in wastewater reuse. However, the latter implies additional post-treatment after the conventional activated sludge processes, in order to produce a safer effluent. Our work aimed at determining the efficiency of reducing the toxicity associated with organic micropollutants (OMPs) in wastewater effluents, using 3 different post-treatment technologies (granular activated carbon (GAC), sand biofiltration and UV irradiation): in particular, target chemical analysis of the OMPs most commonly founded in wastewater was coupled with effect-based assays (estrogenicity and mutagenicity). While chemical analysis assessed satisfactory performances for all 3 technologies in the abatement of selected OMPs, biological assays evidenced another perspective: both GAC and sand biofilters were significantly able to make the estrogenic load plummet; however, the UV system was ineffective in estrogenicity abatement, and its effluent exhibited also a slight mutagenicity, likely due to photo-transformation by-products. These results indicate that a synergistic combination of chemical analysis and biological assays can drive to a proper gauging of post-treatment technologies, taking into account not only the removal of OMPs, but also their overall toxicity.

Keywords: Estrogenicity, Granular activated carbon, Mutagenicity, Organic micropollutants, Sand filtration, UV irradiation

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*Corresponding Author
E-mail: matteo.papa82@gmail.com
Tel: +61 7555 27836 Fax: +61 7555 27836
ORCID: 0000-0003-4872-3411
1. Introduction

The research on organic micropollutants (OMPs) in wastewater has become a worldwide issue of increasing environmental concern, especially considering the growing interest in the reuse of treated wastewater. Thus, the application of post-treatment technologies such as biofiltration systems, UV irradiation or advanced oxidation processes is considered a promising solution to adequate the quality of secondary effluents in view of an eventual reclamation [1].

Despite the large number of studies investigating the fate and removal of various individual and specific groups of OMPs after post-treatments, so far little is known whether these processes affect the toxic activity of OMPs themselves and, consequently, of the effluents. This underdevelopment is mainly due to a lack of standardized protocols for the selection and the execution of biological assays, especially for specific modes of toxic action [2]. Another weakness is related to the use of techniques neither i) specific enough (in reaching the target endpoints) nor ii) adequately sensible. As the first issue is concerned, for instance Vibrio fischeri is frequently employed, but it is only able to detect the baseline toxicity [3]; for the second, the determination of estrogenic activity by means of traditional techniques (e.g. yeast estrogenic screen) does not allow a satisfactory degree of sensibility and robustness, while the application of human breast cancer MCF-7 cells for estrogenicity detection ensures accurate and reproducible results thanks to its high concentration of estrogenic receptors.

Thus, the development of sensible and specific assays represents a valuable strength when profiling tertiary treatments [4, 5]. Indeed, effect-based monitoring approaches are required to gain information on the cumulative effects of complex mixtures characterizing wastewater [6]:

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the detection of OMPs alone is not sufficient to evaluate the ecohazard of treated effluent and, ultimately, the negative effects on aquatic organisms [2, 7, 8].

The objective of this work was to determine the efficiency of three post-treatment technologies (granular activated carbon (GAC) biofiltration, sand biofiltration and UV irradiation) for the mitigation of estrogenic and mutagenic activities (measured by means of MCF-7 cells and Ames test, respectively), therefore expanding the information obtained from the OMPs chemical analysis alone.

2. Experimental Section

2.1. Reactors

2.1.1. Biofiltration systems

Laboratory scale biofilters were used, consisting of two columns of methacrylate, each with 3 L of sand and granular activated carbon as filtering material, respectively. Both biofilters were fed with a secondary effluent (main characteristics: pH: 7.5, chemical oxygen demand (COD): 79 mg/L, ammonium (N-NH$_4$): 0.9 mg/L and nitrate (N-NO$_3$): 5.9 mg/L) that was spiked with selected OMPs (full details in Table S1 of Supplementary Materials). The detailed description of biofilters operation can be found in [9]: among a wider spectrum of operating conditions tested against OMPs removal, the effect-based assays were carried out under an empty bed contact time (EBCT) of 4.8 h. During this period, two sampling campaigns were carried out for quantifying target OMPs and, on the second campaign, bioassays were executed on triplicate and duplicate for estrogenicity and mutagenicity, respectively.
2.1.2. UV system

The UV lab reactor was a 1 L cylindrical glass contactor provided with a low-pressure lamp (TNN 15/32, nominal power 15 W) emitting monochromatic radiation at 254 nm. Similarly to biofilters, a secondary effluent (main characteristics: pH: 7.3, COD: 24 mg/L, N-NH$_4$: 1.3 mg/L, NO$_3$: 3.6 mg/L and turbidity: 4.7 NTU) spiked with selected OMPs was used (full details in Table S1 of Supplementary Materials). The detailed description of UV reactor operation can be found in [10]: again, among a wider spectrum of operating conditions where OMPs were detected, the effect-based assays were carried out under an UV dose of 1,200 mJ/cm$^2$. For this experiment, OMPs were determined on duplicate whereas bioassays were executed on triplicate for estrogenicity and on duplicate for mutagenicity.

It is worth to note that, for both reactors, the treatment conditions we decided to test are stronger than those usually applied in full-scale systems: especially the UV dose, usually in the range of 50 – 200 mJ/cm$^2$ [11]. However, these conditions were properly selected to ensure effectiveness towards not only the parent OMPs, but also their transformation products, which play a key role in the effect-based assays.

2.2. Organic Micropollutants

Several OMPs commonly found in secondary effluents were targeted in this work: three painkillers (ibuprofen, naproxen and diclofenac), four antibiotics (erythromycin, roxithromycin, sulfamethoxazole and trimethoprim), three psychoactive drugs (fluoxetine, carbamazepine and diazepam), three musk fragrances (galaxolide, tonalide and celestolide), three estrogens (estrone, 17β-estradiol and 17α-ethinylestradiol) and four endocrine disruptor compounds - EDCs (4-
octylphenol, 4-nonylphenol, bisphenol A and triclosan). Compound abbreviations are reported in the caption of Fig. 1.

For their determination, 250 mL influent and effluent samples were prefiltered (0.70 µm, Millipore) and pre-concentrated by solid phase extraction (SPE) prior to analysis with Gas Chromatography Mass Spectrometry - GC/MS (pain killers, musk fragrances and EDCs) and Liquid Chromatography Tandem Mass Spectrometry - LC/MS/MS (antibiotics, estrogens and psychoactive drugs). Further details on the limits of detection, quantification and recoveries of analytical methods are described in [9] and [10].

It is worth noting that, as we spiked samples by adding three estrogens in concentration of 1 µg/L (100 – 1,000 times the usual concentration in wastewater), thus E1, E2 and EE2 were by far the major estrogenic active substances, due to a relative estrogenic potency 100 – 1,000 times higher than the other spiked EDCs [12].

2.3. Effect-Based Assays

The same SPE as for chemical analyses was executed, and extracts dried under nitrogen flow and resuspended in 1 mL dimethyl sulfoxide (DMSO), leading thus to an initial enrichment factor of 250 mL<sub>wastewater</sub>/mL<sub>extract</sub>.

2.3.1. Estrogenic activity

Human breast cancer cell line MCF-7 was selected to measure the estrogenic activity. MCF-7 stably transfected with the ERE-tK-LUC construct (kindly supplied by Mikko Unkila, Hormos Medical Ltd, Turku, Finland) was maintained in DMEM (Modified Dulbecco’s Medium, Milan,
Italy), supplemented with 10% fetal bovine serum, at 37°C and 5% CO₂. Cells were plated at a density of 2.5·10⁵ cells/cm² in several plates containing 1 mL of culture medium (phenol red-free DMEM and 5% charcoal-stripped serum). 24 h later, 1 µL of each DMSO extract (therefore, with a dilution factor of 1,000) was added by triplicate and dishes were kept at 37°C for 24 h. As controls without extracts, one cell-plate was supplemented with DMSO solvent, another with ethanol and a last one only with cells. After incubation, cells were harvested and lysed in passive lysis buffer (Promega, Italy). Lysate was spun for 15 s at 12,000 g and supernatant submitted to luciferase activity quantification (Luciferase Assay System, Promega, Italy), by means of a luminometer (GloMAX, Promega, Italy) over 10 s, and expressed as RLU (relative light units) normalized towards protein content (Bradford assay, Biorad, Italy). The latter value was then expressed as estradiol equivalent concentration (ng-EEQ/L bioassay), based on the calibration curve, and ultimately as ng-EEQ/L sample using the relative enrichment factor (REF). REF represents the combination of the initial enrichment factor and the dilution factor in the bioassay plates, as explained by [13]: therefore, REF was equal to 0.25 mL wastewater/mL bioassay.

For the calibration curve, reference estrogen (E2 dissolved in ethanol) was employed, at concentrations corresponding to physiological/sub-physiological doses, i.e. from 10⁻¹⁵ to 10⁻⁸ M. The resulting curve (sigmoidal function) was fitted using Graphpad Prism 6.0 software (GraphPad Software, Inc., USA). Details of cell response to the reference estrogen, together with the calibration curve, are reported in Fig. S1 and Table S2 of Supplementary Materials.
2.3.2. Mutagenic activity

Mutagenicity was assessed using Ames test, which is able to evidence point mutations in bacteria; specifically, we used the TA98 strain of Salmonella typhimurium (able to detect frameshift mutagens) according to [14]. Bacteria were exposed to increasing doses of DMSO extracts: 1-, 5-, 10-, 25- and 50-mL of wastewater equivalents per plate. Positive (10 μg/plate of 2-nitrofluorene) and negative (DMSO solvent) controls were executed.

Data were expressed as mutagenicity ratio (MR), dividing the revertants/plate by the spontaneous mutation rate derived from the negative control. Results were considered positive if MR was higher than 2 (a response at least twice the negative control).

3. Results and Discussion

As aforementioned, we spiked target OMPs: therefore, raw values obtained from bioassays were not discussed themselves, but rather used to calculate and compare removal efficiencies, as hereinafter discussed. Raw data are however reported in Supplementary Materials (Figure S2 for estrogenic and Table S3 for mutagenic activity).

Bioassays were the main focus of this paper, while two previous papers ([9] and [10] that act as accompanying papers) were entirely devoted to the chemical analysis results, specifically explaining the behaviour of OMPs in the three studied post-treatment systems and their different removal mechanisms.
3.1. Biofiltration Systems

First of all, we analyzed chemical data as background information, to see what we should expect from the effect-based assays. Fig. 1 summarizes the removal efficiencies obtained for targeted OMPs: for a quicker understanding, removals were grouped into four categories with appropriate color coding.

It is clear how GAC biofilter exhibited a great potential for OMPs removal: indeed, only the green category (high removal) is represented in Fig. 1, with efficiencies always above 90% for all targeted micropollutants. On the contrary, all the 4 categories are displayed in the case of sand biofilter, where OMPs elimination was strongly related to their biodegradability: for instance, unsatisfactory abatement characterized the most recalcitrant/hardy biodegradable compounds (such as diclofenac, carbamazepine and diazepam); the same compounds, instead, were strongly removed in GAC thanks to the adsorption on activated carbon. Further details of the removal mechanisms in [9].

Then, we examined the bioassays outcomes for the effect-based analysis. Fig. 2 reports the removal of estrogenic activity, calculated on the basis of estradiol equivalent concentrations (ng-EEQ/L, reported in Fig. S2 of Supplementary Materials). Both GAC and sand biofilters reached a strong abatement of estrogenicity (higher than 90%), in agreement with the outcomes of other researches ([15] among others).

To link the removal of the most critical pollutants exhibiting estrogenic activity (i.e., the 3 estrogens) with the reduced estrogenicity, we plotted in Fig. 2 both the information (chemical and biological), and we recorded the accordance of their outcome: indeed, estrogenicity abatement (> 90%) was strongly in agreement with estrogens removal, that also topped 90%
(except EE2 in sand biofiltration, however removed at a 70% extent). This outcome is a confirmation that no estrogenically active by-products were formed during biofiltration, as already postulated by other authors [16]: by using an approach only based on chemical analysis, it would not have been possible to get such a holistic answer, as biotransformation products would not have been taken into account.

On the other hand, mutagenic activity was not observed, neither in the influent nor in the effluent of the biofilters, since the mutagenicity ratio was always below the threshold of 2 (Fig. 3, while the raw values of revertants/plate are reported in Table S3 of Supplementary Materials). This is explained by the fact that spiked OMPs are not recognized as mutagenic compounds, and GAC and sand biofiltration do not generate transformation products able to induce frameshift mutations [16].

Notwithstanding, the lack of transformation products after biofilters is not tout court, but only related to those compounds responsible for monitored biological activities (estrogenicity and mutagenicity): indeed, the formation of transformation products in/during biofiltration is evident elsewhere [17] for other OMPs.

3.2. UV Systems

UV irradiation showed, as also sand biofiltration did, a variety in OMPs behavior, with 3 categories on the ground: some compounds (e.g. diclofenac and estrogens) were remarkably eliminated (green category of Fig. 1), while others (erythromycin, carbamazepine and diazepam) fell in the orange category (i.e. low abatement). As extensively reported in [10], the reason lies
in the different photosensitivity of target OMPs, whose removal efficiencies are very closely related to.

Moreover, it is well known [18] that UV treatment is not able to reach the complete mineralization of pollutants, but only to reduce their complexity owing to the rupture of bonds by UV light action. The transformation of parent compounds into by-products was clearly evidenced by the effect-based estrogenicity assay. Indeed, when we again linked the removal of the most critical pollutants exhibiting estrogenic activity (i.e., the 3 estrogens) with the reduced estrogenicity (Fig. 2), we recorded divergent outcomes: after the UV irradiation the real estrogenic burden measured with the bioassay remained almost unchanged in the effluent (only a 10% removal efficiency), despite the satisfactory abatement of the estrogens (> 80%). As anticipated before, we can therefore attribute this outcome to the formation of transformation products that still possess estrogenic activity, but not detected by the target chemical analysis: for instance, the hydroxylation of estradiol into 2-hydroxyestradiol, which is able to form hydrogen bonds with the estrogen receptor, was observed by [18] after UV. This behavior, therefore, could explain the persistence of the estrogenic activity after UV treatment, as also observed in other works ([19] among others). On the contrary, it is highly unlikely that the low estrogenicity removal could be attributable to the low elimination rate of the UV technique recorded for other micropollutants (CBZ, DZP, etc).

As mutagenicity is concerned, UV system exhibited again some weaknesses (Fig. 4). While the determination of the mutagenicity was not possible in the influent at the highest tested doses (25- and 50-mL wastewater equivalent/plate), because of the strong toxic effect on bacteria, the effluent maintained a borderline mutagenicity strength, with MR values slightly above the
threshold of 2. Similar findings were also achieved by other authors [20], that have shown how UV irradiation could lead to the formation of potential mutagenic by-products.

4. Conclusions

This study represents the last piece of a bigger “puzzle”, preceded by the works [9] and [10] that analysed the behaviour of several organic micropollutants in three post-treatment technologies for secondary effluents polishing. In this last work, their efficiency was evaluated towards two specific modes of toxic actions (the most commonly studied in wastewater: estrogenic and mutagenic activity), with an integrated chemical-biological approach, i.e. by coupling the information derived from OMPs detection with those derived from the effect-based approach (estrogenic and mutagenic activity).

The results indicated that bioassays are needed to correctly gauge the ecohazard posed by micropollutants, as there was no direct link between OMPs removal and toxicity attenuation. In the case of estrogenicity, indeed, although both biofiltration and UV treatment satisfactory eliminated the estrogens (E1, E2, EE2), only the former was able to make the estrogenic load plummet, while UV treatment left it almost unchanged. Moreover, unlike biofilters, after UV treatment also a slight mutagenic effect was recorded. Therefore, effect-based assays suggested how photolysis could generate by-products, missed by the target chemical analysis but still bioactive and therefore detected by bioassays.

In conclusion, the application of a multitiered bio-chemical strategy may allow to properly profile tertiary treatments, and eventually leading to the optimization of their operational conditions in order to obtain a removal not only of OMPs, but also of their toxicity.
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Author Contributions

M.P. (Research Fellow) and L.P. (Research Fellow) designed the experiments, analyzed data and co-wrote the paper; L.P. (Research Fellow) operate the reactors and carried out micropollutants detection; D.F. (Professor) and G.V. (Research Fellow) conducted the mutagenicity tests; G.M. (Professor) and N.S. (Research Fellow) conducted the estrogenicity tests; R.P. (Assistant Professor), J.L. (Professor), F.O. (Professor) and M.C. (Professor) supervised the research and critically reviewed the paper.
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Fig. 1. Average OMPs removal efficiencies in biofilters and UV. Compounds are listed according to their behavior in sand biofilter and labelled as follows: 4-octylphenol (OP), 4-nonylphenol (NP), tonalide (AHTN), galaxolide (HHCB), triclosan (TCS), estrone (E1), ibuprofen (IBP), 17β-estradiol (E2), celestolide (ADBI), erythromycin (ERY), bisphenol A (BPA), roxithromycin (ROX), naproxen (NPX), 17α-ethinylestradiol (EE2), fluoxetine (FLX), sulfamethoxazole (SMX), trimethoprim (TMP), diclofenac (DCF), carbamazepine (CBZ), diazepam (DZP). The colors indicate the level of abatement: green = high (> 75%); yellow = mild (50% to 75%); orange = low (25% to 50%); red = absent (< 25%) (for interpretation of the references to color in this figure legend, the reader is referred to the web version of this article).
**Fig. 2.** Estrogenic bioassay VS chemical analysis. Average removal of estrogenic activity in biofilters and UV (bioassay response), and comparison with chemical analysis (i.e., estrogens removal). The colors indicate the level of abatement: green = high (>75%); yellow = mild (50% to 75%); orange = low (25% to 50%); red = absent (<25%) (for interpretation of the references to color in this figure legend, the reader is referred to the web version of this article).

**Fig. 3.** Average Mutagenicity Ratio obtained using Ames test, for GAC and sand biofilters (n.a. = not available). Background color indicates a negative (green = MR < 2) or positive (red = MR > 2) answer (for interpretation of the references to color in this figure legend, the reader is referred to the web version of this article).
Fig. 4. Mutagenicity ratio obtained using *Ames* test, for UV treatment (tox = toxic to bacteria). Background color indicates a negative (green = MR < 2) or positive (red = MR > 2) answer (for interpretation of the references to color in this figure legend, the reader is referred to the web version of this article).