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# Removal of phenolic compounds in olive mill wastewater by silica–alginate–fungi biocomposites

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**Abstract** This study aims to attempt a treatment strategy based on fungi immobilized on silica-alginate (biocomposites) for removal of phenolic compounds in olive oil mill wastewater (OMW), OMW supplemented (OMWS) with phenolic compounds and water supplemented (WS) with phenolic compounds, thus decreasing its potential impact in the receiving waters. Active (alive) or inactive (death by sterilization) Pleurotus sajor caju was encapsulated in alginate beads. Five beads containing active and inactive fungus were placed in a mold and filled with silica hydrogel (biocomposites). The biocomposites were added to batch reactors containing the OMW, OMWS and WS. The treatment of OMW, OMWS and WS by active and inactive biocomposites was performed throughout 28 days at 25 °C. The efficiency of treatment was evaluated by measuring the removal of targeted organic compounds, chemical oxygen demand (COD) and relative absorbance ratio along the time. Active P. sajor caju biocomposites were able to remove 64.6-88.4 % of phenolic compounds from OMW and OMWS and 91.8-97.5 % in water. Furthermore, in the case of OMW there was also a removal of

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General Physical Chemistry Laboratory, Department of Chemistry, Faculty of Sciences, University Mohammed V-Agdal, Avenue Ibn Battouta, BP 1014, Rabat, Morocco 30.0–38.1 % of fatty acids, 68.7 % of the sterol and 35 % of COD. The silica–alginate–fungi biocomposites showed a high removal of phenolic compounds from OMW and water. Furthermore, in the application of biocomposites to the treatment of OMW it was observed also a decrease on the concentration of fatty acids and sterols as well as a reduction on the COD.

**Keywords** Olive oil mill wastewater · Fungi · *Pleurotus sajor caju* · Biocomposites · Phenolic compounds

## Introduction

Olive mill industry is a seasonal traditional agricultural industry being the producers of the European Union responsible for 74.3 %, Morocco for 5.9 %, Syria for 5.0 %, Tunisia for 5.0 % and Turkey for 4.9 %, among others (International Olive Oil Council 2010). During the seasonal harvesting of olives, a large amount of a liquid waste called olive oil mill wastewater (OMW) is produced and becomes a major environmental problem (Kavvadias et al. 2010; Justino et al. 2012). The OMW constituents depend on several factors such as the soil nature, cultivation system, olive varieties and production process (Roig et al. 2006; McNamara et al. 2008). Some constituents such as tannins, polyalcohols, lipids and phenolic compounds are toxic or recalcitrant, thus rendering the development of alternative treatments to reduce the environmental impact of OMW in natural waters an urgent issue. Many physicochemical treatments, such as ozonation, coagulation, ultrafiltration, electrochemical oxidation, and biological process, using fungi and bacteria were developed and applied for the decolorization, degradation of organic compounds and removal of chemical oxygen demand



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(COD) from OMW (Mantzavinos and Kalogerakis 2005; Borja et al. 2006; Roig et al. 2006; McNamara et al. 2008; Morillo et al. 2009; Cassano et al. 2011; Hanafi et al. 2011; Michailides et al. 2011; Nieto et al. 2011.; Oller et al. 2011; Scoma et al. 2011; Justino et al. 2012). Aerobic biological treatments such as activated sludge appeared to be very effective against some low molecular mass phenolic compounds but are relatively ineffective against the more complex polyphenolics responsible for the dark colouration of OMW (McNamara et al. 2008). In general, fungi have been more effective at degrading both simple phenols and the more complex phenolic compounds present in olivemill wastewaters than bacteria (Morillo et al. 2009), which can be attributed to the structure of the aromatic compounds present in OMW since these compounds are analogous to that of many lignin monomers, and only a few microorganisms, mainly wood-rotting fungi, are capable to efficiently degrade lignin by producing ligninolytic enzymes such as lignin peroxidases, manganese peroxidises and laccases. The treatment of OMW using anaerobic digestion presents several advantages in comparison with classic aerobic process such as the high degree of purification with high-organic-load feeds, low nutrient requirements, low quantities of sludge produced, and generation of biogas (Borja et al. 2006, Morillo et al. 2009). To overcome problems of nutrient imbalance, toxicity and other difficulties derived from the composition of OMW, the wastewaters can be subjected to pre-treatments before anaerobic digestion. The aerobic treatments are traditionally preferred to anaerobic ones (Morillo et al. 2009) in part due to seasonal production of olive oil wastewater.

In the use of aerobic treatments regarding fungi, there is still some constrains arising from the microorganismenvironment interactions such as: (1) growth imbalance caused the introduction of foreign organisms that can be more resilient than the natives; (2) microorganisms metabolism hindered by the presence of large amounts of pollutants (Perullini et al. 2010). These drawbacks can be overcome by the use of immobilized microorganisms in a matrix such as silica since it allows: (1) the protection of cells or microorganisms from harmful environments; (2) the control of their surrounding environment and of their concentration (Kato et al. 2005; Meunier et al. 2010). Therefore, this study aims to attempt a treatment strategy based on the silica-alginate fungi (Pleurotus sajor caju) biocomposites for removal of phenolic compounds in an OMW, thus decreasing its potential impact in the receiving waters and in the environment. To accomplish this objective and in order to ascertain the OMW matrix effect, the treatment strategy was applied to three different samples: OMW, OMW supplemented with phenolic compounds and water supplemented with phenolic compounds.

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## Materials and methods

General characterization of the OMW samples

Samples from OMW were collected from an evaporation pond in the north of Portugal and resulted from a threephase olive oil extraction process made in small scale production facilities. OMW samples were collected in screw-cap glass flasks, transported to laboratory and stored at -20 °C until treatments. OMW samples were characterized for pH ( $4.0 \pm 0.1$ ), COD ( $32.0 \pm 0.6$  g/L), and total phenolic content ( $243 \pm 14$  mg/L). COD was determined following American Society for Testing Materials standard method D 1252-88 (ASTM 1994), pH measured by a pH meter (Crison, Spain), and total phenolic content was determined following the colorimetric method of Folin–Ciocalteau (Mulinacci et al. 2001).

Biological material and culture conditions

Pleurotus sajor caju (São Paulo State University, Brazil) was cultured at 25 °C in a media containing 20 g/L of malt extract, 1 g/L of peptone, and 16 g/L of agar. After the growth period, P. sajor caju was maintained at 4 °C in the culture medium. Before the encapsulation (immobilization procedure), mycelium of P. sajor caju was grown in a liquid medium containing 20 g/L of malt extract and 1 g/L of peptone during 11 days, at 25 °C and 120  $\pm$  10 rpm. After growth, the mycelium was collected by filtering the culture medium through a sterilized gaze, and kept in sterilized plastic containers at 4 °C, until fungus immobilization (Ferreira et al. 2008; Freitas et al. 2009; Justino et al. 2009; Rocha-Santos et al. 2010; Justino et al. 2010). Before the immobilization procedure, the filtered mycelium was processed using a commercial homogenizer (Taurus, Portugal).

## Fungi immobilization

Figure 1 shows the encapsulation procedure used for fungi immobilization, which was adapted from Perullini et al. (2010) and Duarte et al. (2012). First, *P. sajor caju* mycelium was encapsulated in calcium alginate by dropping a suspension of mycelium containing 0.50 % of fungus (dry weight) and 1.25 % (w/v) sodium alginate into a 0.10 M CaCl<sub>2</sub> aqueous solution (needle diameter of 1.2 mm). The alginate beads (Fig. 1A, A1) were collected (after stirring for 10 min) and a total of five beads were placed into each cylindrical glass mould of 1 cm diameter and 1 cm height (Sigma, Portugal). Each mould containing the five beads (Fig. 1B, C) was filled with a mixture of 3 volumes of sodium silicate (0.83 M in Si(IV) (Sigma, Portugal) and 1 volume of commercial colloidal silica Ludox HS40 (Sigma, Portugal), adjusting pH to 4.5 (hydrochloric acid and stirring for 30 s). A nanoporous monolithic structure (Fig. 1D) was obtained by the silicate polymerization in the presence of silica nanoparticles. The monoliths were kept at room temperature over night in a clean chamber until their introduction in the batch reactor. In order to infer about the contribution of both fungi and monolithic structure, moulds containing the monolithic structure with inactive encapsulated fungi were also prepared for comparison purposes. To inactivate the fungi, the mycelium was sterilized at 121 °C, during 16 min before encapsulation.

Fungi alginate beads and filled molds with fungi-alginate beads were observed by optic microscope (OM) through  $40 \times$  objective lenses and images were photographed using a digital camera. Images of the surface of the monoliths were obtained with a scanning electronic microscope (SEM).

## OMW treatment

Figure 2 displays a schematic view of the OMW treatment. The monolithic structures with encapsulated active (A) and inactive (I) *P. sajor caju* (biocomposites) were added to 40 mL of OMW and to OMW supplemented (OMWS) with phenolic compounds (24 mg/L of p-coumaric acid, 22 mgL of vanillin, 20 mg/L of caffeic acid, 26 mg/L of vanillic acid, or 20 mg/L of tyrosol) in batch reactors (two mould per batch reactor) (Justino et al. 2010). A 25 % (v/v) of OMW was used for the treatment since the efficiency of biological treatment depends on composition and dilution

of the OMW (Ferreira et al. 2008; Justino et al. 2009). Dilutions were made with deionised water and the following additives were added to the diluted OMW: 1.0 g/L of potassium dihydrogen phosphate, 0.405 g/L of diammonium tartrate dibasic, and 0.05 g/L of yeast extract. pH was adjusted to  $4.02 \pm 0.05$  before starting treatment (Ferreira et al. 2008; Justino et al. 2009).

In order to ascertain if there is a matrix effect in the degradation of phenols in OMW, the essay performed with OMW and OMWS was totally repeated with deionised water supplemented with phenolic compounds (WS).

The batch reactors with 40 mL of OMW or OMW supplemented with phenolic compounds (OMWS) or water supplemented with phenolic compounds (WS) with active biocomposites (AB) or inactive biocomposites (IB) (Fig. 2A, A1) were kept at 25 °C and stirring at 120  $\pm$  10 rpm, during 28 days of treatment. A total of 50 samples were used for each category (OMW, OMWS and WS). During this period, samples corresponding to AB and IB treatments were collected after 0, 7, 14, 21 and 28 days, for OMW, OMWS and WS samples. The treated samples were then analyzed for relative absorbance ratio and COD. The samples after 28 days of treatment were also evaluated for phenolic and other organic compounds previously targeted by Justino et al. (2009).

## Liquid-liquid extraction

Samples (OMW, OMWS and WS) resulted from 0 and 28 days of treatment (40 mL per replicate) were submitted to liquid–liquid extraction (Justino et al. 2010). Therefore,

Fig. 1 Procedure for immobilization of fungi in alginate-silica: A microphotographs from OM of fungi-alginate bead amplified 400 times; A1 photograph of fungi-alginate beads with 2 mm Ø; B microphotographs from OM of five fungi-alginate beads into the mold filled with silica amplified 400 times; C photograph of the 1 cm Ømold with five fungi-alginate beads; D microphotograph from SEM of the surface of the monoliths (500 µm)





Fig. 2 Treatment scheme of OMW. A Microphotographs from OM of five fungi–alginate beads into the mold filled with silica amplified 400 times; A1 photograph of the mold with five fungi–alginate beads before treatment; B microphotographs from OM of five fungi–alginate beads into the mold filled with silica after 7 days of treatment amplified 400 times; B1 photograph of the mold with five fungi–alginate beads after 28 days of treatment



40 mL of each sample and 25 mL of diethyl ether were mixed in a separation funnel, collecting by aspiration the upper layer. Extraction was repeated twice with 25 mL of diethyl ether and 25 mL of acetate ethyl, respectively. The organic extracts were then filtered through sodium sulfate, collected in 100 mL pear-shaped flask and placed in a rotavapor at 45 °C until a volume of 0.5 mL. The 0.5 mL extract was kept under N<sub>2</sub> until dryness. Dried samples were re-dissolved with 3 mL diethyl ether and transferred to a micro vial. Sample derivatization was performed by addition of 250  $\mu$ L of pyridine (Fluka, Spain), 250  $\mu$ L of bis(trimethylsilyl)trifluoracetamide (Fluka, Spain), and 50  $\mu$ L of chlorotrimethylsilane (Sigma-Aldrich, Germany) and then, holding the mixture at 70 °C, in a sand bath for 30 min.

# GC-MS analysis

For quantification of phenolic and other organic compounds, the extracts from samples OMW, OMWS and WS resulted from 0 and 28 days of treatment were analyzed by GC–MS (Shimadzu QP 5000, Japan) equipped with a capillary column (SPB-5; 30 m  $\times$  0.32 mm; 0.25 µm film thickness; Supelco, Spain). The following



oven temperature program was used: 80-220 °C at 3 °C/min, then 220–290 °C at 6 °C/min, and maintained at 290 °C for 3.09 min. The carrier gas used was helium with an inlet pressure of 50 kPa (1 min) to 90 kPa (1.5 kPa/min) and then to 110 kPa (0.7 kPa/min). The GC–MS was operated in selected ion monitoring mode to obtain the quantification of organic compounds. Table 1 shows the parameters for measurement ions (Mz).

The concentration of each compound was determined by direct interpolation in the standard curve within their linear dynamic range, and the detection limits (Table 1) were calculated using  $y = y_B + 3s_B$ , where  $s_B$  is the SD of the blank signal estimated as  $s_{y/x}$ , the residual SD taken from the calibration line, and  $y_B$  is the blank signal estimated from the intercept taken also from the calibration line (Miller and Miller 2005).

## Absorbance ratio

Absorbance scan (200–600 nm) and fixed measures at 270 nm wavelength of the samples collected at different treatment periods were obtained by UV–VIS spectro-photometry (GBC/Cintra 10e). Before absorbance measurements, the pH was adjusted to 4.0 by the addition of

Table 1 Parameters for measurement ions and detection limits

Compound	Measurement parameters (MZ)	Detection limit (ng/L)		
Vanillin	224	0.25		
Tyrosol	282	0.32		
Caffeic acid	194	0.40		
Vanillic acid	182	0.56		
p-coumaric acid	178	0.43		
Myristoleic acid	240	0.75		
Palmitic acid	270	0.67		
Linoleic acid	294	0.12		
Oleic acid	296	0.49		
Stearic acid	298	0.98		
α-Linolenic acid	292	0.53		
Lignoceric acid	382	0.76		
β-Sitosterol	414	0.15		

sodium hydroxide (0.1–1 M) or hydrochloric acid (0.1–1 M), and samples were filtered through GF/F (47 Ø mm, Whatman) glass-fiber filters and diluted 7.5 times with deionised water (pH 4.0). The absorbance at 270 nm was measured since it is attributed to  $\pi$ – $\pi$ \* electronic transitions in phenolic compounds and phenolic acids.

## Quality assurance and quality control

In order to assure representativeness and reliability of the results obtained, QC/QA procedures included blanks, duplicates and recovery assays. Recovery assays were performed to ensure the representativeness of the extraction procedure of phenolic compounds from OMW samples. Replicate analysis of the samples before and after treatments gave an uncertainty of <5% for COD and organic compounds.

## Statistical analysis

All statistical analyses were performed using SigmaStat<sup>TM</sup> 3.0 (1994). One-way ANOVA was carried out to verify if there was a statistically significant difference for the removal of organic compounds between treatments with active and inactive biocomposites for each sample and also between OMWS, OMW and WS samples. Before the application of one-way ANOVA, normality and homoscedasticity of experimental data were verified through the Kolmogorov–Smirnov test, to verify if independent data variances were normally distributed, and the Levene test to evaluate the equality of variances, respectively (SigmaStat<sup>TM</sup> 3.0 1994).

## **Results and discussion**

Table 2 displays the concentration of organic compounds in the OMW. It can be observed that the concentration of phenolic compounds (phenols and phenolic acids) ranged from 6.3 µg/L for vanillic acid to 20.2 µg/L for caffeic acid whereas the concentration of fatty acids ranged from 6.1 µg/L for lignoceric acid to 88.7 µg/L for oleic acid. It can also be observed in Table 3 that the recovery values for the three replicate samples of OMW were about 100 % for both the two phenols and three phenolic acids added to the effluent. The concentration of organic compounds present in this OMW was different from the hydroxytyrosol in OMW as reported by De Marco et al. (2007). This fact could be related with several factors such as the extraction process, olives, olive ripening, olive storage duration before milling and storage of the effluent in evaporation ponds. The recovery values evidence a high extraction efficiency of the liquid-liquid procedure for the extraction of phenols and phenolic acids from OMW.

Table 3 shows the removal percentages of organic compounds in OMW, OMWS and WS, respectively. The removal percentages of organic compounds on OMW and OMWS by both active and inactive biocomposites treatment throughout time did not show a statistically significant difference (P = 0.971 for AB treatment; P = 0.989 for IB treatment), indicating that the extra concentration of phenolic compounds added to OMW was not a limiting factor for the treatment by both active or inactive biocomposites.

There was a statistical significant difference on the percentage of removal of organic compounds achieved for the treatments by active and inactive biocomposites (OMW: P = 0.001, WS:  $P \le 0.001$ ). In the OMW, the higher values of removal percentages were observed for phenols (88.4 and 33.5 % for tyrosol by AB and IB treatments, respectively), while the lowest percentages were achieved for fatty acids (30.0 % for  $\alpha$ -linolenic acid by AB treatment and 26 % for lignoceric acid by IB treatment). Concerning the WS in comparison with OMW and OMWS, higher values of removal percentage of phenols and phenolic acids were achieved by AB and IB treatments. As an example of success in the removal of phenols by laccase (a enzyme generally expressed by *P. sajor caju*) immobilized on sol-gel-derived silica, Qiu and Huang (2010) used chlorophenols as model for phenolic compounds and achieved a removal percentage between 90 and 100 % for 2,4-dichlorophenol and 2,4,6-trichlorophenol in sole solutions of deionized water. However, when Qiu and Huang (2010) mixed both phenols in the solution, the removal rates of both compounds declined. Therefore, in our work, the differences between the removal of phenolic compounds in OMW, OMWS and in water (WS) could be



Table 2 Concentration of organic compounds in OMW

Compound (µg/L)	OMW	Supplemented (mg/L)	Recovery	
Phenols				
Vanillin	$16.2\pm0.6$	22.00	100.1	
Tyrosol	$18.3\pm0.5$	20.00	100.4	
Phenolic acids				
Caffeic acid	$20.2\pm0.9$	20.00	100.2	
Vanillic acid	$6.3\pm0.4$	26.00	99.8	
p-coumaric acid	$19.0\pm0.8$	24.00	99.9	
Fatty acids				
Myristoleic acid	$8.5\pm0.9$	-	-	
Palmitic acid	$25.5\pm0.7$		-	
Linoleic acid	$26.3\pm0.9$	-	-	
Oleic acid	$88.7\pm0.8$	-	-	
Stearic acid	$11.2\pm0.3$	-	-	
$\alpha$ -Linolenic acid	$6.5\pm0.5$	-	-	
Lignoceric acid	$6.1\pm0.7$	-	-	
Sterols				
β-Sitosterol	$12.3\pm0.6$	-	100.1	

attributed to matrix effect namely to the presence of polymeric compounds in OMW makes the treatment more difficult (McNamara et al. 2008). The average composition of OMW is 83-89 % water, 3.5-15 % of organic matter and 0.5-2 % of mineral salts (Asses et al. 2009). About 10 % of the organic matter of the OMW has been related with polymeric phenolic compounds that display a lignin-like structure, which constitutes the most recalcitrant fraction (Justino et al. 2012). In fact, the treatment by active biocomposites can be considered as a three-step process responsible for the removal of the compounds (Chen et al. 2005; Svec and Huber 2006): (1) adsorption of reactants on the monolithic structure and diffusion to the biological active sites; (2) biodegradation by the fungi, and (3) diffusion of the products resulting from the biodegradation. Typically, low molecular weight compounds diffuse relatively quickly through the monolith structure reaching easily to the immobilized fungi, whereas the transfer of high molecular weight compounds is considerably slower due naturally to lower diffusion coefficients.

In the treatment by inactive biocomposites (IB), only the adsorption/diffusion phenomena are expected since the metabolic pathways of P. sajor caju were inactivated by sterilization. Therefore, the significant differences (OMW: P = 0.001, WS:  $P \le 0.001$ ) achieved in the removal of organic compounds (e.g. Tyrosol 88.4 and 33.5 %) by active and inactive biocomposites can be attributed to the biodegradation performed by P. sajor caju and therefore by the enzymes from this fungi (Chen et al. 2005; Svec and



Table 3 Removal percentages of organic compounds (phenols, phenolic acids, fatty acids and sterol) in OMW, OMWS and WS after 28 days of treatments by active biocomposites (AB) and inactive biocomposites (IB)

Compound	Removal (%)						
	OMW	OMW		OMWS		WS	
	AB	IB	AB	IB	AB	IB	
Vanillin	78.9	34.6	78.9	34.6	93.2	81.6	
Tyrosol	88.4	33.5	88.4	33.5	97.5	85.3	
Caffeic acid	72.1	33.8	72.1	33.8	92.4	80.6	
Vanillic acid	64.6	30.1	64.6	30.1	91.8	79.9	
p-coumaric acid	74.8	32.9	74.8	32.9	96.0	84.8	
Myristoleic acid	31.8	27.2	31.8	27.2	_	-	
Palmitic acid	31.1	28.0	31.1	28.0	_	-	
Linoleic acid	35.2	26.2	35.2	26.2	_	-	
Oleic acid	37.2	29.4	37.2	29.4	_	-	
Stearic acid	34.3	29.5	34.3	29.5	_	_	
α-Linolenic acid	30.0	26.8	30.0	26.8	_	_	
Lignoceric acid	33.5	26.0	33.5	26.0	_	_	
β-Sitosterol	68.7	31.1	68.7	31.1	_	-	

Huber 2006), since they are present only in the active biocomposite.

Figure 3 shows the absorbance ratio [Abs(t)/Abs(t0)] at 270 nm throughout time of treatment of OMW, OMWS and WS by active and inactive biocomposites, respectively.

It can be observed in Fig. 3 that the absorbance decreases in all the samples until 7 days of treatment, remaining thereafter approximately constant until 28 days of treatment. Symbols and lines in Fig. 3 almost overlap for OMWS and OMW during treatments by active (OMWS-AB and OMW-AB) biocomposites since there was no statistically significant difference between them (P = 0.976). The same behavior was observed for both samples during treatments by inactive (OMWS-IB and OMW-IB) biocomposites (P = 0.983). The absorbance ratio in OMW and OMWS decreased 25 and 15 % after 28 days of treatments by active and inactive biocomposites, respectively. In turn, the absorbance ratio of WS decreased 94 and 82 % due to the treatments by active and inactive biocomposites, respectively. The observed differences between the values achieved in WS and OMW could be attributed to the presence of polymeric phenols (ligninlike compounds) in the composition of OMW, as discussed previously for the removal of phenolic compounds. In fact, the 25 % of absorbance ratio decrease (Fig. 3) in OMW compared to the 65-89 % of removal of the phenolic compounds (Table 3) promoted by active biocomposites treatment reinforces the idea that the lignin-like macromolecular compounds (McNamara et al. 2008) were not removed, making treatment of OMW difficult.



**Fig. 3** Absorbance ratio [Abs(*t*)/Abs(*t*0)] at 270 nm throughout time of treatment. OMWS-IB, OMW supplemented with phenolic compounds treated by inactive biocomposites; OMWS-AB, OMW supplemented with phenolic compounds treated by active biocomposites; OMW-IB, OMW treated by inactive biocomposites; OMW-AB, OMW treated by active biocomposites; WS-IB, water supplemented with phenolic compounds and treated by inactive biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites; WS-AB, water supplemented with phenolic composites; WS-AB, water supplemented wi

Figure 4 shows the percentage of removal of COD in OMW, OMWS and WS throughout time of treatments. Symbols and lines in Fig. 4 almost overlap for OMWS and OMW during treatments by active (OMWS-AB and OMW-AB) biocomposites since there was no statistically significant difference between them (P = 0.981). The same behavior was observed for both samples during treatments by inactive (OMWS-IB and OMW-IB) biocomposites (P = 0.973). The percentage of removal of COD in all samples increased until 7 days, reaching thereafter a plateau until the end of treatments, which is in accordance with the variation of absorbance ratio between the last 21 days of treatment. The highest removal of COD after 28 days of treatments was achieved in WS treated by active biocomposites (92 %) and inactive biocomposites (83 %), while the lowest removal of COD was attained in both OMW and OMWS treated by active (35 %) and inactive biocomposites (30 %). These data reinforce once more that not all organic compounds were removed.

After 28 days of treatment of OMW and OMWS, the active biocomposites were removed from the effluent and introduced in untreated effluent following the procedure described in 2.4 for OMW treatment in order to verify their capability to re-utilization. After another 28 days of treatment, there was no statistically significant difference between the removal of phenolic compounds (P = 0.979) by the active biocomposites used by the first time or re-used after 28 days (data not shown).



Fig. 4 Percentage removal of COD (%) throughout time of treatment. OMWS-IB, OMW supplemented with phenolic compounds treated by inactive biocomposites; OMWS-AB, OMW supplemented with phenolic compounds treated by active biocomposites; OMW-IB, OMW treated by inactive biocomposites; OMW-AB, OMW treated by active biocomposites; WS-IB, water supplemented with phenolic compounds and treated by inactive biocomposites; WS-AB, water supplemented with phenolic compounds and treated by active biocomposites

# Conclusion

The treatment of OMW by biocomposites showed to be a potential alternative way to not only decrease the concentration of phenolic compounds but also to decrease the concentration of fatty acids and sterols as well as to reduce COD. This treatment also showed its potential to remove phenolic compounds in water.

Some limitations to degrade high molecular weight compounds are, however, a drawback that limits the full application of biocomposites to treat effluents. Further research combining a previous degradation of high molecular weight compounds could be a possible way to optimize the treatment of OMW by biocomposites.

The treatments with biocomposites using immobilized fungi are positive aspects that should be considered from environmental point of view since it allows to control and prevent the release of fungi into the environment.

The biocomposites showed ability to be re-used and therefore this treatment can be interesting from an economic point of view.

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