

# Diclofenac and 2-anilinophenylacetate degradation by combined activity of biogenic manganese oxides and silver

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## Summary

The occurrence of a range of recalcitrant organic micropollutants in our aquatic environment has led to the development of various tertiary wastewater treatment methods. In this study, biogenic manganese oxides (Bio-MnOx), biogenic silver nanoparticles (Bio-Ag<sup>0</sup>) and ionic silver were used for the oxidative removal of the frequently encountered drug diclofenac and its dechlorinated form, 2-anilinophenylacetate (APA). Diclofenac was rapidly degraded during ongoing manganese oxidation by *Pseudomonas putida* MnB6. Furthermore, whereas preoxidized Bio-MnOx, Bio-Ag<sup>0</sup> and Ag<sup>+</sup> separately did not show any removal capacity for diclofenac, an enhanced removal occurred when Bio-MnOx and silver species were combined. Similar results were obtained for APA. Finally, a slow removal of diclofenac but more rapid APA degradation was observed when silver was added to manganese-free *P. putida* biomass. Combining these results, three mechanisms of diclofenac and APA removal could be distinguished: (i) a co-metabolic removal during active Mn<sup>2+</sup> oxidation by *P. putida*; (ii) a synergistic interaction between preoxidized Bio-MnOx and silver species; and (iii) a (bio)chemical process by biomass enriched with silver catalysts. This paper demonstrates the use of *P. putida* for water treatment purposes and is the first report of the application of

silver combined with biogenic manganese for the removal of organic water contaminants.

## Introduction

Since the discovery of a range of pharmaceuticals and personal care products in surface water, groundwater and even drinking water (Ternes, 1998; Heberer, 2002; Mompelat *et al.*, 2009), tertiary treatment methods were developed to remove micropollutants after conventional wastewater treatment. Physicochemical techniques for the removal of micropollutants are mostly based upon filtration and oxidative processes (Ikehata *et al.*, 2006) but may have specific disadvantages, such as high costs (Johnson and Sumpter, 2001), the formation of toxic by-products (Guzzella *et al.*, 2002; Richardson *et al.*, 2007) and inefficiency against certain compounds (Ternes *et al.*, 2003). Biogenic metals have shown to be useful catalysts for the removal of recalcitrant organic compounds. Biogenic palladium nanoparticles are efficient catalysts for the reductive transformation of halogenated organics, such as diatrizoate (Hennebel *et al.*, 2010; De Gusseme *et al.*, 2011a) and diclofenac (De Gusseme *et al.*, 2012). However, these reductive treatments are generally limited to a dehalogenation step, upon which other treatments are required to remove the formed reaction products. As such, the end-product of a reduction of diclofenac with biogenic palladium is its dechlorinated form 2-anilinophenylacetate (APA) (De Corte *et al.*, 2011; De Gusseme *et al.*, 2012). Conversely, oxidative biological techniques of micropollutant removal are under investigation. For example, biogenic manganese(III,IV) oxides (Bio-MnOx) can be successfully used for the removal of a range of pharmaceuticals (Forrez *et al.*, 2010; 2011). The oxidative power of manganese oxide compares with that of oxygen gas (Thauer *et al.*, 1977). In addition, the efficiency of micropollutant removal is further improved by the presence of manganese-oxidizing bacterial biomass: a continuous cyclic reoxidation of manganese can theoretically result in a sustained removal of micropollutants (de Rudder *et al.*, 2004; De Schamphelaire *et al.*, 2007) and oxidation products might be further degraded by the bacterial metabolism, increasing the driving force of the initial reaction. Finally, evidence exists that during the oxidation of Mn<sup>2+</sup> to Mn<sup>4+</sup> by bacteria such as *Pseudomonas putida*

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and *Bacillus* sp. SG-1, ligand-bound  $Mn^{3+}$  intermediates are produced that might sensitively increase the oxidative power of a manganese-bacteria mixture (Kostka *et al.*, 1995; Parker *et al.*, 2004; Murray and Tebo, 2006).

In water treatment, the disinfecting activity of both chemically and biologically produced silver nanoparticles (Bio-Ag<sup>0</sup>) has been shown valuable for the inactivation of bacteria and viruses (Jain and Pradeep, 2005; De Gussemme *et al.*, 2011b). Additionally, silver nanoparticles can be used as catalysts in a range of organic redox reactions (Jiang *et al.*, 2005; Xu *et al.*, 2006). Whereas silver compounds are generally weak redox catalysts, studies have demonstrated a sensitive increase in activity for composite metal catalysts. For example, Haruta and Sano (1983) demonstrated a higher activity of Ag–Co and Ag–Mn composite oxides, in comparison with the single-component oxides, for the oxidation of hydrogen gas to H<sub>2</sub>O and carbon monoxide to CO<sub>2</sub>. Comparable results were obtained for the catalytic oxidation of volatile organic compounds, such as acetone and pyridine (Luo *et al.*, 1998).

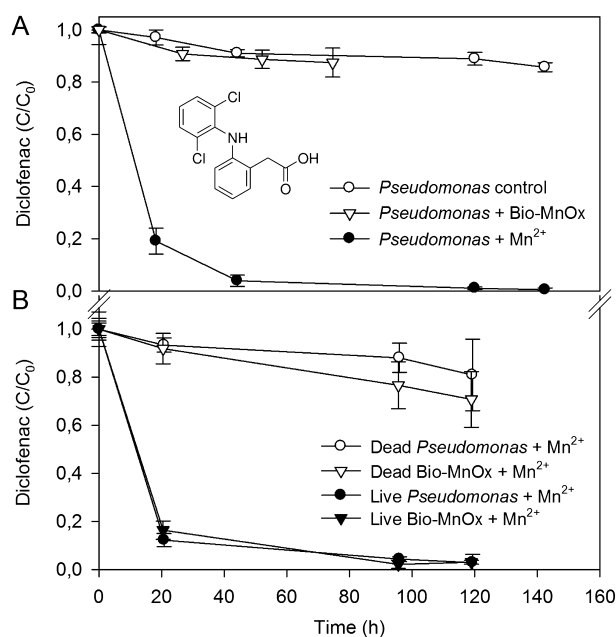
This study has explored the applicability of Bio-MnOx and silver species for the oxidative removal of micropollutants during water treatment, using the chlorinated pharmaceutical compound diclofenac as a model pollutant. The main objectives were (i) to evaluate the influence of the manganese-oxidizing metabolism of *P. putida* on the removal of diclofenac with Bio-MnOx; (ii) to determine the effects of an enrichment of manganese oxides with silver species on the degradation of diclofenac; and (iii) to assess the degradation of 2-anilinophenylacetate, the dechlorinated product of diclofenac.

## Results

### Influence of manganese-oxidizing metabolism on diclofenac removal

Diclofenac (3 mg l<sup>-1</sup>) and manganese chloride (3.28 mg Mn<sup>2+</sup> l<sup>-1</sup>) were added simultaneously to a *P. putida* culture. In a parallel setup, diclofenac was added 113 h after the addition of Mn<sup>2+</sup>, to allow a complete oxidation of manganese by the bacteria prior to the addition of diclofenac. Diclofenac was readily removed in batches where manganese was simultaneously oxidized, with a decrease of 96 ± 2% within 44 h (first-order rate constant  $k = 0.065 \text{ h}^{-1}$ ), while a lower decrease of 13 ± 6% after 75 h ( $k = 0.0023 \text{ h}^{-1}$ ) was observed in batches where manganese was oxidized prior to diclofenac addition (Fig. 1A). Manganese-free control experiments, in which an equal amount of bacteria was used, showed a diclofenac removal of 14 ± 2% after 142 h.

To address the importance of bacterial metabolism during manganese oxidation and thus diclofenac degradation, control experiments using heat inactivated cul-



**Fig. 1.** Removal of diclofenac by preformed Bio-MnOx and by active manganese-oxidizing metabolism of *P. putida*. A. During oxidation of 3.28 mg Mn<sup>2+</sup> l<sup>-1</sup> (*Pseudomonas* + Mn<sup>2+</sup>), after complete oxidation of Mn<sup>2+</sup> (*Pseudomonas* + Bio-MnOx) and in manganese-free controls (*Pseudomonas* control). B. In heat-inactivated *P. putida* cultures supplied with 3.28 mg Mn<sup>2+</sup> l<sup>-1</sup> (‘dead *Pseudomonas* + Mn<sup>2+</sup>’), in heat-inactivated Bio-MnOx (3.28 mg Mn l<sup>-1</sup>) supplied with another 1.64 mg Mn<sup>2+</sup> l<sup>-1</sup> added (‘dead Bio-MnOx + Mn<sup>2+</sup>’), and in living controls. C<sub>0</sub> = 3 mg l<sup>-1</sup> diclofenac. Error bars indicate standard deviations of three replicates; sometimes smaller than symbols.

tures of *P. putida* were set up (Fig. 1B). Upon addition of 3.28 mg Mn<sup>2+</sup> l<sup>-1</sup> and 3 mg l<sup>-1</sup> diclofenac to non-viable biomass, no formation of biogenic manganese oxides could be observed and a slight removal of 19 ± 15% diclofenac was seen after 119 h of incubation. In contrast, after the addition of equal amounts of manganese and diclofenac to a viable *P. putida* culture, repeating the active manganese oxidation of the previous experiment shown in Fig. 1A, a faster removal of diclofenac was observed (97 ± 3% after 119 h, Fig. 1B).

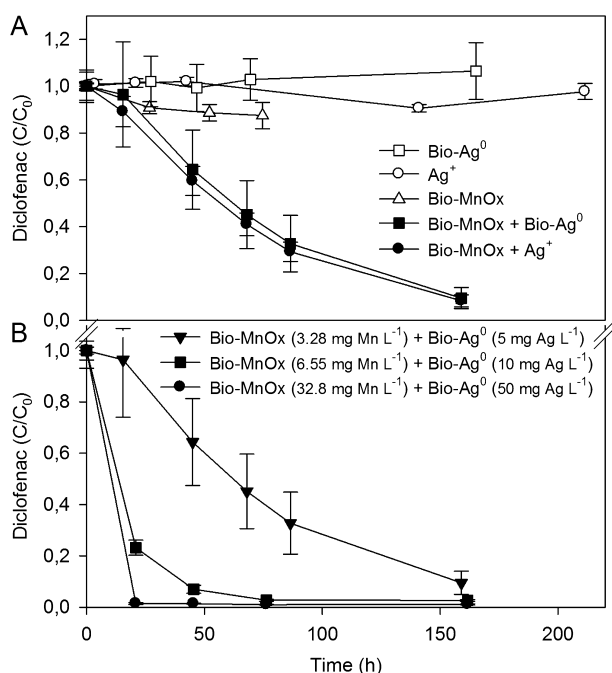
Subsequently, it was tested whether preformed Bio-MnOx could facilitate the reoxidation of Mn<sup>2+</sup> in an autocatalytic manner, rendering the bacterial manganese-oxidizing metabolism unnecessary for the ensuing removal of diclofenac. To this extent, heat inactivation of preformed Bio-MnOx (3.28 mg Mn l<sup>-1</sup>) was performed, after which an additional amount of 1.64 mg Mn<sup>2+</sup> l<sup>-1</sup> and 3 mg l<sup>-1</sup> diclofenac were added. The diclofenac concentration decreased only to a limited extent (29 ± 12% after 119 h), which was in contrast to the diclofenac removal after addition of equal amounts of Mn<sup>2+</sup> and diclofenac to viable Bio-MnOx. Indeed, the latter amounted to 97% ± 1 after 119 h (Fig. 1B), which was again comparable to the

diclofenac removal rate during active manganese oxidation obtained in the previous experiments.

#### Improved diclofenac removal in the presence of silver

To further improve the reactivity of manganese oxides, the influence of silver species on the degradation of diclofenac by Bio-MnOx was assessed. When diclofenac was added to a mixture of preformed Bio-MnOx (3.28 mg Mn l<sup>-1</sup>) and either ionic silver or biogenic nanoparticles (Bio-Ag<sup>0</sup>) (5 mg Ag l<sup>-1</sup>), removal percentages of 92 ± 3% and 91 ± 5%, respectively, were obtained after 159 h (first-order rate constants  $k = 0.017$  and  $0.016$  h<sup>-1</sup>). Control experiments containing Bio-Ag<sup>0</sup>, Ag<sup>+</sup> (5 mg Ag l<sup>-1</sup>) or Bio-MnOx (3.28 mg Mn l<sup>-1</sup>) separately showed little to no diclofenac removal (Fig. 2A) ( $k = 0$  h<sup>-1</sup>,  $0.0003$  h<sup>-1</sup> and  $0.0023$  h<sup>-1</sup> respectively).

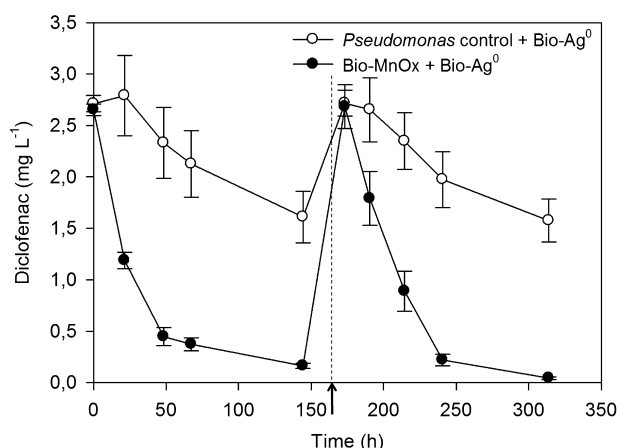
At increased biomass and metal concentrations, diclofenac removal rates were sensitively enhanced (Fig. 2B). When increasing the concentrations of biomass, manganese and silver by a factor 2 and 10, a better removal of 97 ± 0.4% after 76 h and 99 ± 0.4% after 21 h, respectively, could be obtained. This corre-



**Fig. 2.** Removal of diclofenac by combined manganese and silver species.

A. By Bio-MnOx (3.28 mg Mn l<sup>-1</sup>) combined with Bio-Ag<sup>0</sup> or Ag<sup>+</sup> (5 mg Ag l<sup>-1</sup>), and by Bio-Ag<sup>0</sup>, Ag<sup>+</sup> and Bio-MnOx as separate controls.

B. Diclofenac removal at elevated metal concentrations: 3.28 mg Mn l<sup>-1</sup> as Bio-MnOx with 5 mg l<sup>-1</sup> silver as Bio-Ag<sup>0</sup>, and twofold and 10-fold increases. C<sub>0</sub> = 3 mg l<sup>-1</sup> diclofenac. Error bars indicate standard deviations of three replicates; sometimes smaller than symbols.



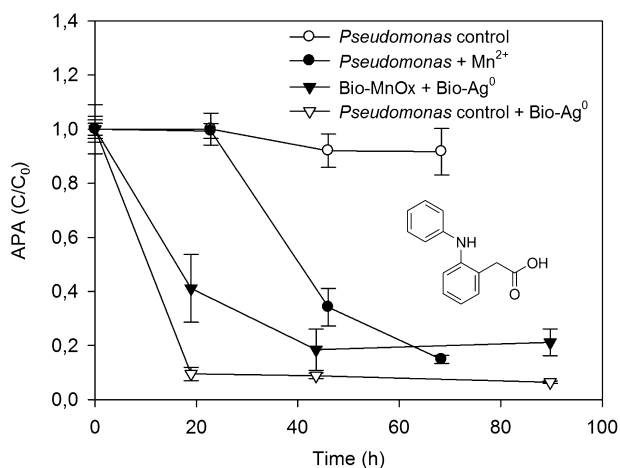
**Fig. 3.** Removal of diclofenac by Bio-Ag<sup>0</sup> (10 mg Ag l<sup>-1</sup>) combined with *P. putida* biomass and with preoxidized Bio-MnOx (6.55 mg Mn l<sup>-1</sup>). C<sub>0</sub> = 3 mg l<sup>-1</sup> diclofenac. The arrow marks a respiration of diclofenac to 3 mg l<sup>-1</sup> at time  $t = 173$  h. Error bars indicate standard deviations of three replicates; sometimes smaller than symbols.

sponded with a factor 3.6 and 5.7 increase in the diclofenac removal rate constants ( $k = 0.058$  h<sup>-1</sup> and  $0.091$  h<sup>-1</sup> respectively).

To distinguish degradation processes from sorption, a respiration experiment was performed for diclofenac removal by Bio-MnOx (6.55 mg Mn l<sup>-1</sup>) and Bio-Ag<sup>0</sup> (10 mg Ag l<sup>-1</sup>) (Fig. 3). Diclofenac was removed by 94 ± 1% after 144 h during a first incubation period. After 173 h, the batches were respiked to the initial concentration of 3 mg l<sup>-1</sup> diclofenac. During a second incubation period, diclofenac was removed at a comparable rate, with a decrease of 98 ± 0.5% after 141 h. A parallel control experiment, containing manganese-free *P. putida* biomass with Bio-Ag<sup>0</sup>, showed a diclofenac removal by 41 ± 9% after 144 h during the first incubation and 42 ± 7% after 141 h during the second incubation period.

#### Removal of APA

To examine the structural importance of the chlorine moieties in the diclofenac molecule and their contribution to its degradability by manganese and silver species, degradation experiments were performed using 2-anilinophenylacetate (APA) (Fig. 4). When 3.28 mg Mn<sup>2+</sup> l<sup>-1</sup> and 3 mg APA l<sup>-1</sup> were added to a viable *P. putida* culture, APA was removed by 85 ± 2% after 45 h during active manganese oxidation, after a lag phase of 23 h in which no visible formation of Bio-MnOx or APA removal was observed. In contrast, control experiments using manganese-free biomass showed no APA removal. In the presence of preoxidized Bio-MnOx (3.28 mg Mn l<sup>-1</sup>) and Bio-Ag<sup>0</sup> (5 mg Ag l<sup>-1</sup>), APA was removed by 82 ± 8% after 44 h. Surprisingly, a control experiment containing a



**Fig. 4.** Removal of APA by *P. putida* during active oxidation of Mn<sup>2+</sup> (3.28 mg l<sup>-1</sup>); by preoxidized Bio-MnOx (3.28 mg Mn l<sup>-1</sup>) combined with Bio-Ag<sup>0</sup> (5 mg Ag l<sup>-1</sup>); and by manganese-free controls. C<sub>0</sub> = 3 mg APA l<sup>-1</sup>. Error bars indicate standard deviations of three replicates; sometimes smaller than symbols.

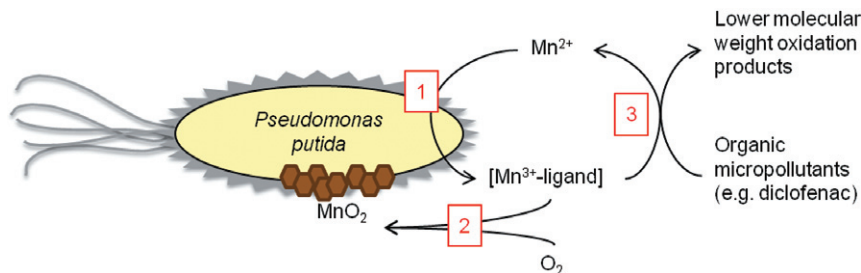
mixture of Bio-Ag<sup>0</sup> (5 mg Ag l<sup>-1</sup>) and a manganese-free *P. putida* culture showed an even faster removal, with a decrease of APA concentration by 90 ± 2% after 19 h.

## Discussion

Traditional wastewater treatment is focused on the removal of the organic load (chemical oxygen demand, COD) and nutrient concentrations before discharge in natural aquatic systems. However, with continuously increasing global water demands, the need to develop techniques for wastewater reuse arises. Of specific concern for the reuse of water is the presence of persistent organic compounds, such as certain pharmaceuticals, pesticides and industrial chemicals. This research has elaborated on the use of biogenic manganese oxides as novel agents for the removal of micropollutants, in continuation of the research of Sabirova and colleagues (2008) and Forrez and colleagues (2010; 2011). Moreover, this study used for the first time silver as a catalyst, combined with manganese oxides, for the removal of micropollutants.

## Removal of diclofenac during active manganese oxidation

Almost no removal of diclofenac in the presence of preformed Bio-MnOx was observed. In contrast, during active Mn<sup>2+</sup> oxidation by *P. putida*, diclofenac was readily removed, with almost a 30-fold increase of the first-order rate constant (Fig. 1). This suggests a participating role of the manganese-oxidizing metabolism of the bacteria in the removal of diclofenac, especially since heat-inactivated cultures showed no considerable diclofenac removal. Although it is known that the autocatalytic oxidation of Mn<sup>2+</sup> may occur on the surface of preformed manganese oxides (Murray, 1975), these results show that possible intermediates during the autocatalytic oxidation of MnO<sub>2</sub> do not contribute to the degradation of diclofenac, but that, in contrast, the metabolic activity of the biomass is the most significant factor herein. The oxidation of manganese by *P. putida* likely involves a multicopper enzyme, which oxidizes Mn<sup>2+</sup> to a ligand-bound Mn<sup>3+</sup> intermediate (Parker *et al.*, 2004; Tebo *et al.*, 2004; Webb *et al.*, 2005). This highly reactive Mn<sup>3+</sup> intermediate is then thought to be further oxidized to Mn(IV) oxide by putative enzymes in the extracellular mucous layer of the bacteria (De Vrind *et al.*, 2003). A three-step model of the oxidizing activity of Bio-MnOx can be proposed (Fig. 5): (i) Mn<sup>2+</sup> is oxidized to a ligand-bound Mn<sup>3+</sup> intermediate, presumably by a multicopper enzyme of *P. putida*; (ii) this Mn<sup>3+</sup> intermediate can either be further oxidized to Mn(IV) oxide, plausibly by enzymes on the cell surface; or (iii) the reactive Mn<sup>3+</sup> intermediate can oxidize other compounds, such as micropollutants, upon which Mn<sup>2+</sup> is formed again. As such, the degradation of diclofenac can be considered a co-metabolic process during the oxidation of manganese. The model presented here is in concordance with Forrez and colleagues (2010), who observed that diclofenac removal by Bio-MnOx was inhibited when *P. putida* was treated with sodium azide, a biostatic agent. Further study should focus on the technical challenge of verifying this model by demonstrating the formation of unstable Mn<sup>3+</sup> intermediates and correlating their presence with the oxidation of diclofenac. Moreover, research on mutant strains should allow to identify the different



**Fig. 5.** Proposed model of the oxidation of Mn<sup>2+</sup> by *P. putida* and the co-metabolic degradation of micropollutants such as diclofenac: (1) oxidation of Mn<sup>2+</sup> to a ligand-bound Mn<sup>3+</sup> intermediate (after Parker *et al.*, 2004; Tebo *et al.*, 2004; Webb *et al.*, 2005); (2) oxidation of Mn<sup>3+</sup> to MnO<sub>2</sub> (after De Vrind *et al.*, 2003); (3) reduction of Mn<sup>3+</sup> to Mn<sup>2+</sup> with concomitant oxidation of the micropollutant.

*P. putida* enzymes involved in the oxidation of manganese and clarify the mechanism of diclofenac oxidation during this process.

#### Removal of diclofenac by the activity of silver species

To increase the overall reactivity,  $\text{Ag}^+$  and  $\text{Bio-Ag}^0$  were added to  $\text{Bio-MnOx}$ , upon which diclofenac was rapidly removed. A respirometry experiment (Fig. 3) clearly refuted the possibility of sorption as the prime mechanism of diclofenac removal by this silver –  $\text{Bio-MnOx}$  mixture, suggesting a catalytic nature of the process. Silver and biogenic manganese oxides cooperate in a synergistic manner, increasing the diclofenac removal rate to a higher level than would be expected from combining the removal rates of both separately:  $\text{Ag}^+$ ,  $\text{Bio-Ag}^0$  and  $\text{Bio-MnOx}$  alone showed little to no diclofenac removal (Fig. 2). Likewise, Imamura and colleagues (1988) observed similar synergistic interactions between manganese and silver oxides for the oxidation of gaseous carbon compounds.

A surprising increase of the diclofenac removal rate was already observed when  $\text{Bio-Ag}^0$  was combined with manganese-free biomass (Fig. 3). One possible explanation could involve the formation of reactive oxygen species (ROS), such as hydroxyl radicals, superoxide and peroxide, since both ionic silver (Matsumura *et al.*, 2003; Park *et al.*, 2009) and silver nanoparticles (Su *et al.*, 2009; Sintubin *et al.*, 2011) have been shown to interact with the cell wall and intracellular respiratory enzymes of bacteria, thereby creating ROS. Given the susceptibility of diclofenac towards oxidation by ROS (Hofmann *et al.*, 2007; Sein *et al.*, 2008), it is considered plausible that the observed removal of diclofenac in the presence of  $\text{Bio-Ag}^0$  in a *P. putida* culture occurs through this reaction.

In this study, no definite conclusions could be made regarding the mechanism of the observed  $\text{Bio-MnOx}$  – silver interaction during the degradation of diclofenac. Yet, the results hitherto suggest a twofold mode of action: (i) silver species ( $\text{Ag}^+$  and  $\text{Bio-Ag}^0$ ) interact with *P. putida* biomass in such a way that diclofenac is slowly removed; and (ii) silver species enhance the reactivity of  $\text{Bio-MnOx}$ , in such a way that diclofenac is more rapidly removed. It is suggested that further research should attempt to demonstrate an increased amount of ROS when the bacterial cells are triggered by silver and the subsequent effects on micropollutants such as diclofenac. Likewise, the mechanism of the observed catalytic manganese – silver synergy should be addressed. Furthermore, given the antibacterial activity of silver (Morones *et al.*, 2005; Bjarnsholt *et al.*, 2007; Yoon *et al.*, 2007), careful study should determine whether silver may be added to active *P. putida* biomass at concentrations sufficiently low not to inhibit the oxidation of manganese and the associated diclofenac removal, while still allowing the catalytic

silver –  $\text{Bio-MnOx}$  interaction to further enhance diclofenac removal rates.

#### Differences in reactivity between diclofenac and APA

The end-product of a reductive treatment of diclofenac with, for example, palladium or palladium – gold catalysts is 2-anilinophenylacetate (APA) (De Corte *et al.*, 2011; De Gussemme *et al.*, 2012). In order to explore the use of manganese and silver species to complement such reductive treatments, additional removal experiments were conducted with APA. During active bacterial oxidation of manganese, APA was rapidly removed after a 23-hour lag phase, in which no manganese oxide formation and APA removal occurred (Fig. 4). After this delayed onset of manganese oxidation, the APA removal rate was comparable to that of diclofenac under equal circumstances. Remarkably, when silver was added to a manganese-free *P. putida* culture, a faster APA removal was observed than when combined with  $\text{Bio-MnOx}$ . Thus, it is concluded that silver acts as an efficient catalyst for the removal of APA in the presence of *P. putida* biomass and that manganese is not required in this reaction.

A comparison of the degradation kinetics of diclofenac with those of APA might provide more insight in the structural importance of the chlorine moieties of diclofenac and in the mechanism by which it is transformed. The main bacterial degradation pathway of diclofenac is an oxidation to 5-hydroxydiclofenac, followed by (auto)oxidation to diclofenac-2,5-iminoquinone (Gröning *et al.*, 2007). Likewise, these two compounds were the only metabolites detected for diclofenac degradation by  $\text{Bio-MnOx}$  (Forrez *et al.*, 2010), although they only accounted for 5–10% of the total diclofenac removal. In this study, both diclofenac and APA were removed at comparable rates during the ongoing oxidation of manganese by *P. putida*. Therefore, it is concluded that the chlorine moieties of diclofenac do not affect its principal degradation pathway, which is likely to involve a hydroxylation of the ring structure. When added to a combination of  $\text{Bio-MnOx}$  and  $\text{Bio-Ag}^0$ , APA was removed at a lower rate than in a manganese-free bacterial culture containing only  $\text{Bio-Ag}^0$  (Fig. 4), suggesting that manganese oxides are not required for the degradation of APA by  $\text{Bio-Ag}^0$  and even interfere with it. On the contrary, an enhanced removal rate was observed for diclofenac in the presence of both manganese oxides and silver, compared with silver in a manganese-free bacterial culture (Fig. 3). It is therefore concluded that the principal degradation mechanism of diclofenac by manganese and silver might differ from that of APA and that the chlorine moieties of diclofenac counter its degradability by silver in manganese-free biomass. Possibly, APA is, more than diclofenac, sensitive to degradation by the ROS that are produced extra- and intracellularly in the presence of

silver ions. This might explain the higher APA removal rate in the absence of Bio-MnOx, since deposits of manganese oxides on the bacterial cell wall are expected to obstruct cross-membrane transport. Diclofenac, on the other hand, is primarily degraded as the result of a hitherto unspecified synergistic interaction between manganese oxides and silver.

#### Conclusive remarks

This research has shown that biogenic manganese and silver species can be used as novel agents for the oxidative and catalytic removal of micropollutants. Further research should focus on determining the range of different micropollutants that can be removed by these agents and optimizing the removal conditions, as well as unravelling the reaction mechanisms and identifying the formed metabolites. Additionally, careful study should allow assessing and improving the economic feasibility of water treatment with Bio-MnOx and Bio-Ag<sup>0</sup>, in order to transform these promising, innovative materials into an effective solution to the problem of recalcitrant organic water pollution.

## Experimental procedures

### Bacterial growth conditions

Pure cultures of *P. putida* MnB6 (BCCM/LMG 2322) were grown on a platform shaker (111 r.p.m.) at 28°C under aerobic conditions, in 100 ml or 250 ml Erlenmeyer flasks. The growth medium was prepared according to Boogerd and de Vrind (1987). The cultures were incubated until an optical density (610 nm) of  $1.0 \pm 0.3$  was reached, measured after syringe-and-needle homogenization.

### Diclofenac and APA removal during active manganese oxidation

*Pseudomonas putida* cultures with optical density (610 nm) of  $1.0 \pm 0.3$  were enriched with manganese chloride (3.28 mg Mn<sup>2+</sup> l<sup>-1</sup>) from a stock solution of 500 mg MnCl<sub>2</sub> l<sup>-1</sup> (218 mg Mn<sup>2+</sup> l<sup>-1</sup>) that was previously filter sterilized (0.22 µm, Millipore) and stored at 4°C. Diclofenac was added at the desired time from a cool-stored stock solution in distilled water. Likewise, APA was added from a cool-stored stock solution in 70% ethanol. Experiments were incubated in 100 ml or 250 ml Erlenmeyer flasks, leaving a headspace volume equal to or greater than the volume of the culture liquid, to ensure aerobic conditions. For heat-inactivation experiments, Erlenmeyer flasks containing *P. putida* biomass or preoxidized Bio-MnOx were placed in hot water for 10 min at 60°C. All experiments were performed in triplicate. Between sampling points, batches were stored on a platform shaker (111 r.p.m.) at 28°C. Manganese-free *P. putida* controls were treated identically.

### Diclofenac and APA removal by preformed Bio-MnOx and silver species

Preoxidized Bio-MnOx was produced according to Forrez and colleagues (2010). In brief, *P. putida* cultures were grown to an optical density (610 nm) of  $1.0 \pm 0.3$  and enriched with manganese chloride (3.28 mg Mn<sup>2+</sup> l<sup>-1</sup>) as described above, and stored on a platform shaker at 28°C during 24 h for a complete oxidation of the Mn<sup>2+</sup>. The Bio-MnOx were then centrifuged (10 min at 7500 g) and washed twice (10 min at 10 000 g) with phosphate buffer (10 mM KH<sub>2</sub>PO<sub>4</sub>/K<sub>2</sub>HPO<sub>4</sub>, pH 7–8). The pellets were resuspended at the desired concentration in a test solution containing 0.6 g NaCl l<sup>-1</sup> and 10 mM phosphate buffer (pH 7). Manganese-free control cultures were treated identically.

The production of Bio-Ag<sup>0</sup> was performed following the protocol of Sintubin and colleagues (2009): *Lactobacillus fermentum* G2/10 was grown in Man-Rogosa-Sharpe (MRS) broth (Oxoid) at 28°C on a platform shaker (111 r.p.m.) for 48 h. The biomass was harvested and washed twice with Milli-Q water by means of centrifugation (5 min at 10 000 g). After determination of the cell dry weight (CDW ≈ cell wet weight × 0.23), the biomass was resuspended in Milli-Q water to a concentration of 4.6 g CDW l<sup>-1</sup> and alkalinized with 0.025 M NaOH. Silver (1 g l<sup>-1</sup>) was added in the diamine-complexed form, from a stock solution of silver nitrate that was dosed with ammonium hydroxide until all formed silver precipitates were redissolved. After 24h of incubation (28°C, 111 r.p.m.), the biomass and the formed Bio-Ag<sup>0</sup> were washed thrice (15 min, 10 000 g) with Milli-Q water. The concentration of silver, present on the biomass as Bio-Ag<sup>0</sup>, was determined by atomic absorption spectroscopy (AA-6300 atomic absorption spectroscope, Shimadzu) after addition of 2 ml HNO<sub>3</sub> (65%) and 2 ml H<sub>2</sub>O<sub>2</sub> (30%) to a 2 ml Bio-Ag<sup>0</sup> sample and subsequent heating until the solution was clear.

According to the experimental setup, Bio-Ag<sup>0</sup> and Ag<sup>+</sup>, from a freshly made stock solution of AgNO<sub>3</sub> in Milli-Q water, were added to the Bio-MnOx solution or to Milli-Q water in 100 ml Erlenmeyer flasks. After addition of diclofenac or APA (3 mg l<sup>-1</sup>), batches were stored on a platform shaker (111 r.p.m.) at 28°C between sampling points. All experiments were performed in triplicate.

### Analytical procedures

Samples for diclofenac and APA measurements were filtered with regenerated cellulose filters (RC, 0.20 µm, Phenex) or polyvinylidene fluoride filters (PVDF, 0.22 µm, Millipore), which both tested negative for sorption of diclofenac and APA (data not shown). Prior to analysis, the samples were stored in glass vials at 4°C. Measurements were made using a Dionex P580 HPLC system with a TCC-100 column oven and a diode array detector (UVD340U) and processing was done using Chromeleon 6.80 software. A Genesis C18 column (150 × 4.6 mm, 4 µm, Alltech) with a guard column was used for separation. The injection volume was 50 µl. The separation method was modified from Forrez and colleagues (2010): at a temperature of 40°C and a flow rate of 1 ml min<sup>-1</sup>, the elution was isocratic with 80% HPLC grade methanol and 20% of a solution of 0.1% formic acid in Milli-Q water. Diclofenac was detected by absorbance at a wavelength of 203 nm, at a retention time of 4 min. APA was analysed by

HPLC as described for diclofenac. The absorbance was measured at a wavelength of 275 nm and APA appeared at a retention time of 3 min. Standards ranging from 0 to 5 mg l<sup>-1</sup> were prepared for calibration. The limit of detection (LOD), i.e. the lowest non-zero point in the calibration series, was 100 µg l<sup>-1</sup>.

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