Performance of the Electro-Oxidation and Electro-Fenton Processes with a BDD Anode for the Treatment of Low Contents of Pharmaceuticals in a Real Water Matrix

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Performance of the Electro-Oxidation and Electro-Fenton Processes with a BDD Anode for the Treatment of Low Contents of Pharmaceuticals in a Real Water Matrix

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Abstract: Here, we report the performance of electro-oxidation and electro-Fenton with a boron-doped diamond (BDD) anode for the treatment of single and multicomponent solutions containing small amounts of pharmaceutical residues (i.e., 1 mg·L⁻¹ paracetamol and/or 1 mg·L⁻¹ salicylic acid) spiked into a real water matrix at pH 3.0. Electro-oxidation was performed in a BDD/Pt cell, whereas electro-Fenton was carried out in a BDD/air-diffusion cell to electrogenerate H₂O₂ at the cathode, always operating at constant current density. It was found that the decay of both pharmaceuticals by electro-oxidation was more rapid in the real water matrix than in ultrapure water with 0.05 mol·L⁻¹ Na₂SO₄ because of their additional reaction with active chlorine species produced at the bulk from the oxidation of Cl⁻ ion. Such chlorinated oxidants exhibited even higher reactivity than hydroxyl radicals formed and confined at the anode. The increase in current density largely enhanced the removal of both pollutants. Similar results were found using the real water matrix at natural alkaline pH. When the mixture of both pharmaceuticals was treated by electro-oxidation, their abatement became slower owing to the competitive attack of generated oxidants over them. Only a slight acceleration of pharmaceutical decay was obtained for the real water matrix using electro-Fenton, since the accumulation of additional homogeneous hydroxyl radical formed from Fenton’s reaction between generated H₂O₂ and added Fe²⁺ was inhibited by its reaction with Cl⁻ to form much less reactive chlorinated radicals. For the real water matrix with added pharmaceuticals, a high degree of mineralization of the natural organic matter content (NOM) was reached at high current densities by electro-oxidation, which was even improved upon addition of 0.05 mol·L⁻¹ Na₂SO₄. Traces of oxidation by-products like p-benzoquinone, as well as NO₃⁻ and NH₄⁺ ions, were detected during the electro-oxidation of paracetamol solutions, but the N-compounds contained in the real water matrix were not removed under the investigated conditions.

Key words: boron-doped diamond anode; electro-oxidation; electro-Fenton; paracetamol; real water matrix; salicylic acid

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Over the last fifteen years, pharmaceutical drugs have received increasing attention as potential bioactive chemicals in the environment[1]. They are considered as emerging pollutants in waterbodies because they remain unregulated or their directives and legal frameworks are not yet set-up[2]. Pharmaceuticals and their metabolites are continuously introduced in the aquatic environment, where they are detected at trace concentrations (i.e., as micropollutants found in the ng·L⁻¹ or μg·L⁻¹ range). This pollution arises from emission from production sites, direct disposal of overplus drugs in households and hospitals, excretion

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from urine or faeces after drug administration to humans and animals, and water treatments in fish farms[1]. Pharmaceuticals persist in the environment mainly because of their incomplete elimination in sewage treatment plants (STPs), with 60% to 90% of the parent molecules still present after the action of biodegradation, deconjugation, partitioning, and photodegradation steps[6]. This affects the water quality and drinking water supply, and may constitute a potential risk for the ecosystems, the human and animal welfare in the long term[5]. Although the effects of pharmaceuticals on living beings are not well documented[1], several studies have shown that they may cause slow but irreversible changes to the genome sequence of microorganisms, which increases their resistance to them[2,3]. Also, some drugs classified as endocrine disrupting compounds (EDCs) cause harmful effects on the human endocrine system[2,4,5]. To avoid their potential toxicity and other possible dangerous health effects, the removal of pharmaceuticals and their metabolites from waters must be ensured by developing more powerful oxidation methods.

Recently, a large variety of electrochemical advanced oxidation processes (EAOPs) have been developed to mineralize persistent and toxic organics found in waters[5,7-9]. EAOPs are able to electrogenerate the hydroxyl radical (OH), the second strongest oxidant known after fluorine. It has such a high standard reduction potential (E(\text{OH}/H_2O) = 2.80 V/SHE at 25 °C) that it can non-selectively react with most organics giving dehydrogenated or hydroxylated derivatives, which can eventually be completely mineralized to CO_2, water and inorganic ions[7-9]. The simplest and most common EAOP for water remediation is electro-oxidation, in which organics contained in a contaminated solution are oxidized by direct charge transfer at the anode (M), or rather destroyed by physisorbed hydroxyl radical (M(\text{OH})) formed as intermediate of O_2 evolution from water oxidation at high current densities[7-10]:

\[ \text{M} + \text{H}_2\text{O} \rightarrow \text{M(\text{OH})} + \text{H}^+ + \text{e} \]  (1)

The best anode for electro-oxidation is the boron-doped diamond (BDD) thin-film electrode, which possesses technologically important properties like an inert surface with low adsorption properties, remarkable corrosion stability even in strongly acidic media, and extremely high O_2-evolution overvoltage. These characteristics enhance the removal of organics with reactive BDD(\text{OH})[7], and confer to the BDD anode a high ability to mineralize aromatic pollutants[3,7,8,43] and their by-products, i.e., aliphatic carboxylic acids[24-25], showing much higher oxidation power than traditional anodes like Pt[12] and PbO_2[16].

EAOPs based on H_2O_2 generation have also received great attention for water treatment[9]. The most popular of these methods is electro-Fenton, which involves the continuous supply of H_2O_2 to an acidic contaminated solution from the two-electron reduction of O_2 gas at a carbonaceous cathode, usually carbon felt[17-22] or carbon-polytetrafluoroethylene (PTFE) gas (O_2 or air) diffusion electrodes[17,18,23-25]:

\[ \text{O}_2 + 2\text{H}^+ + 2 \text{e} \rightarrow \text{H}_2\text{O}_2 \]  (2)

In electro-Fenton, a catalytic amount of Fe^{3+} is added to the solution to react with H_2O_2 giving homogeneous *OH and Fe^{2+} from Fenton’s reaction (3). This reaction is catalytic and is mainly propagated by the cathodic reduction of Fe^{3+} to Fe^{2+} from Reaction (4)[3,8].

\[ \text{Fe}^{3+} + \text{H}_2\text{O}_2 \rightarrow \text{Fe}^{2+} + \text{OH} + \text{OH}^- \]  (3)

\[ \text{Fe}^{2+} + \text{e} \rightarrow \text{Fe}^{2+} \]  (4)

Thus, the electro-Fenton process in an undivided cell with a BDD anode involves the attack of organics by both, heterogeneous BDD(\text{OH}) formed from Reaction (1) and homogeneous \text{OH} produced in the bulk from Fenton’s reaction (3).

In previous works, we have applied the electro-oxidation and electro-Fenton processes to effectively mineralize high contents of several pharmaceuticals or metabolites, including triclosan[17], triclocarban[17], chlorophene[18], clofibric acid[23], ibuprofen[26], paracetamol (N-(4-hydroxyphenyl)acetamide)[11,25-26] and salicylic acid (2-hydroxybenzoic acid)[27] in synthetic water matrices. However, much less is known about the viability of these processes for the treatment of low concentrations of pharmaceuticals in surface and ground waters. These aqueous effluents contain natural organic matter (NOM), which is all the
organic matter in a reservoir or natural ecosystem other than living organisms and man-made compounds, so it includes a wide variety of dissolved complex molecules like humic, fulvic, and tannic acids, as well as polysaccharides, among other substances. This dissolved fraction of NOM is hardly oxidized, not being fully removed upon application of conventional water processes in STPs. Lately, more efficient AOPs have emerged for the removal of NOM, aiming to avoid many serious problems related to its presence when water treatments are applied.

To clarify the influence of NOM on the oxidation ability of the above EAOPs, we have undertaken a study on the removal of paracetamol and salicylic acid spiked into a real water matrix. Paracetamol, a common analgesic and anti-inflammatory for humans and animals, has been found with concentrations <10 μg·L⁻¹ in European and Australian STP effluents, as well as in USA natural waters. Salicylic acid, easily produced from hydrolytic deacetylation of the drug acetylsalicylic acid (aspirin), is accumulated in STP effluents and natural waters. Other authors have also shown that paracetamol and salicylic acid can be degraded by both, electro-oxidation with different anodes and electro-Fenton in synthetic aqueous matrices.

This paper reports the performance of two EAOPs, electro-oxidation and electro-Fenton with a BDD anode, for the treatment of 1 mg·L⁻¹ of paracetamol and salicylic acid, alone or mixed, upon addition to a real water matrix. Comparative experiments with ultrapure water were made to study the influence of NOM on the processes. Electro-oxidation was tested with BDD/Pt and BDD/air-diffusion cells to investigate the influence of electrogenerated H₂O₂ on the degradation ability. The effects of current density (j) and Na₂SO₄ content on electro-oxidation performance were also examined.

1 Materials and Methods

1.1 Chemicals

Paracetamol was of reagent grade from Sigma-Aldrich. Salicylic acid, as well as ferrous sulfate heptahydrate used as a catalyst, was of analytical grade from Fluka. Anhydrous sodium sulfate, used as background electrolyte, and sulfuric acid, used to adjust the initial pH of solutions, were of analytical grade from Merck. Synthetic solutions were prepared with ultrapure water obtained from a Millipore Milli-Q system with resistivity > 18 MΩ·cm at 25 °C. Organic solvents and other chemicals used were either of HPLC or analytical grade supplied by Aldrich, Merck, and Panreac.

1.2 Electrolytic Systems

All the electrolytic experiments were conducted in an open, undivided cylindrical cell of 150 mL capacity, equipped with a double jacket where external thermostated water was recirculated to maintain the solution temperature at 25 °C. Solutions of 100 mL with 1 mg·L⁻¹ paracetamol and/or salicylic acid were electrolyzed under vigorous stirring with a magnetic bar at 400 rpm to ensure mixing and the transport of reactants towards/from the electrodes. The anode was always a BDD electrode purchased from Adamant Technologies, synthesized by the hot filament chemical vapor deposition technique on single-crystal p-type Si(100) wafers (0.1 Ω·cm, Siltronix). The cathode was either a Pt sheet purchased from SEMP-SA, used in the majority of electro-oxidation experiments, or a carbon-PTFE air-diffusion electrode supplied by E-TEK for electro-Fenton as well as for some electro-oxidation experiments. This air-diffusion cathode was mounted as described elsewhere and was fed with air pumped at 300 mL·min⁻¹ to generate H₂O₂ from Reaction (2). The geometric area of all the electrodes was 3 cm², and the separation between the anode and cathode in both BDD/Pt and BDD/air-diffusion cells was close to 1 cm. The assays were performed at constant j of 10 or 50 mA·cm⁻², using an Amel 2053 potentiostat-galvanostat as a current source. The potential difference of the cell was directly measured on a Demestres 653B digital multimeter. In order to remove the impurities of the BDD surface and activate the air-diffusion cathode, they were previously polarized in 0.05 mol·L⁻¹ Na₂SO₄ solution at 100 mA·cm⁻² for 60 min. Electro-Fenton treatments were made for solutions of pH 3.0 after
addition of 0.15 mmol·L⁻¹ Fe²⁺. Such values of solution pH and Fe²⁺ concentration were chosen because they were found to be the optimal ones for similar degradation studies of high concentrations of paracetamol[29] and salicylic acid[30] in synthetic solutions.

1.3 Instruments and Analytical Procedures

The solution pH was measured with a Crison GLP 22 pH-meter. Samples were withdrawn at regular time periods from treated solutions and microfiltered with 0.45 µm PTFE filters (Whatman) before analysis. Total carbon was determined with a Shimadzu VCSN TOC analyzer. This system was also used for the mineralization of solutions from their dissolved organic carbon (DOC) decay measured with the non-purgeable organic carbon (NPOC) method. Reproducible DOC values with ±1% precision were found by injecting 50 µL aliquots into the TOC analyzer. Total nitrogen (TN) was obtained with a Shimadzu TNM-1 unit coupled to this analyzer.

Paracetamol and salicylic acid removals were assessed by reversed-phase HPLC using a Waters 600 LC fitted with a Thermo BDS Hypersil C18 5 µm, 250 mm x 4.6 mm, column at 35 °C, and coupled to a Waters 996 photodiode array detector selected at the λ_{max} of the UV spectra of each compound. These analyses were made with a 30:70 (V/V) acetonitrile/water (phosphate buffer of pH 3.25) mixture at 0.6 mL·min⁻¹ as mobile phase, showing well-defined peaks for paracetamol at retention time (t_r) = 5.8 min and λ = 246 nm, salicylic acid at t_r = 8.8 min and λ = 234 nm, and p-benzoquinone at t_r = 7.9 min and λ = 244 nm.

Metal cations (Mg²⁺, Ca²⁺, and iron ions) were quantified by inductively coupled plasma-optical emission spectrometry (ICP-OES) with a Perkin Elmer Optima 3200RL system. Other inorganic ions were detected by ion chromatography using a Shimadzu 10 Avp HPLC coupled to a Shimadzu CDD 10 Avp conductivity detector. Concentrations of Na⁺, K⁺, and NH₄⁺ were obtained by using a Shodex IC YK-421, 125 mm x 4.6 mm, cation column under circulation of a mobile phase of 5.0 mmol·L⁻¹ tartaric acid, 2.0 mmol·L⁻¹ dipicolinic acid, 24.2 mmol·L⁻¹ boric acid, and 15.0 mmol·L⁻¹ crown ether at 1.0 mL·min⁻¹ and 40 °C. Contents of Cl⁻, ClO₃⁻, NO₃⁻, and SO₄²⁻ were determined using a Shim-Pack IC-A1S, 100 mm x 4.6 mm, anion column and a 2.4 mmol·L⁻¹ tris(hydroxymethyl)aminomethane, and 2.5 mmol·L⁻¹ phthalic acid solution of pH 4.0 at 1.5 mL·min⁻¹ and 40 °C as mobile phase.

1.4 Real Water Matrix

The raw water matrix was collected from the primary decantation effluent of a municipal STP in Manresa (Barcelona, Spain) and conserved in a refrigerator at 4 °C before the degradation assays. The characteristics of this real water are summarized in Tab. 1. As can be seen, the total carbon was very high compared to DOC (which was due to dissolved NOM alone), thus suggesting the presence of large amounts of HCO₃⁻ and CO₃²⁻ ions in the raw water at such an alkaline pH. Its conductance was very low compared to that of a 0.05 mol·L⁻¹ Na₂SO₄ solution (10.5 mS), because of the small content of inorganic salts. No ClO₃⁻, NO₃⁻, NH₄⁺, and iron ions were detected, whereas relatively high concentrations of Cl⁻ were found, which can presumably affect the oxidation ability of the EAOPs under study[35,70].

Tab. 1 Main physicochemical characteristics of the real water matrix obtained from a municipal STP

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Total carbon/(mg·L⁻¹)</td>
<td>180</td>
</tr>
<tr>
<td>DOC/(mg·L⁻¹)</td>
<td>8.0</td>
</tr>
<tr>
<td>Total nitrogen/(mg·L⁻¹)</td>
<td>9.0</td>
</tr>
<tr>
<td>Conductance, G (mS)</td>
<td>1.7</td>
</tr>
<tr>
<td>pH</td>
<td>7.9</td>
</tr>
<tr>
<td>[Na⁺]/(mg·L⁻¹)</td>
<td>159</td>
</tr>
<tr>
<td>[K⁺]/(mg·L⁻¹)</td>
<td>52</td>
</tr>
<tr>
<td>[Ca⁺]/(mg·L⁻¹)</td>
<td>111</td>
</tr>
<tr>
<td>[Mg²⁺]/(mg·L⁻¹)</td>
<td>24</td>
</tr>
<tr>
<td>[Iron ions]/(mg·L⁻¹)</td>
<td>Not detected</td>
</tr>
<tr>
<td>[Cl⁻]/(mg·L⁻¹)</td>
<td>273</td>
</tr>
<tr>
<td>[SO₄²⁻]/(mg·L⁻¹)</td>
<td>189</td>
</tr>
</tbody>
</table>
2 Results and Discussion

2.1 Electro-Oxidation of Solutions Containing Paracetamol Alone

Fig. 1 shows the comparative decay of 1 mg·L⁻¹ of paracetamol from both, synthetic solutions with 0.05 mol·L⁻¹ Na₂SO₄ and the real water matrix, at different pH values using a BDD/Pt cell at 50 mA·cm⁻².

As can be seen, this pharmaceutical is completely removed in synthetic solutions by electro-oxidation with a BDD anode, in 45 min at pH 3.0 and more slowly, in 60 min, at natural pH (ca. 6.0). These results indicate the production of slightly higher quantities of oxidant BDD(∙OH) from Reaction (1) in the former medium, in agreement with the electrochemical characterization of BDD reported elsewhere [39]. It should be noted that in these trials the solution pH did not vary significantly during the electrolysis times tested. On the other hand, Fig. 1 also shows that paracetamol decay is strongly enhanced using the real water matrix, being removed at a similar rate at both, natural pH (7.9) and pH 3.0, with total disappearance in only 5 min. The large enhancement in the oxidation rate of this pharmaceutical in a real water matrix can be contributed by the generation of active chlorine species from the parallel oxidation of the Cl⁻ ion (see Tab. 1) at the BDD anode. Therefore, such chlorinated oxidants and BDD(∙OH) become competitors.

It is well known [7, 40-41] that Cl⁻ ion is directly oxidized at the BDD anode to yield soluble chlorine by Reaction (5), which diffuses away from the anode to be rapidly hydrolyzed and disproportionated to hypochlorous acid and chloride ion from Reaction (6). Hypochlorous acid is then in equilibrium with hypochlorite ion from Reaction (7) with pKₐ = 7.55.

\[
\begin{align*}
2\text{Cl}^- & \rightarrow \text{Cl}_2(aq) + 2e^- \quad (5) \\
\text{Cl}_2(aq) + \text{H}_2\text{O} & \rightarrow \text{HClO} + \text{Cl}^- + \text{H}^+ \quad (6) \\
\text{HClO} & \rightarrow \text{ClO}^- + \text{H}^+ \quad (7)
\end{align*}
\]

From the above reactions, it can be concluded that the predominant chlorine species in the bulk are Cl₂ up to pH = 3, HClO in pH = 3 ~ 8, and ClO⁻ at pH > 8 [42].

It is thus expected that the mediated oxidation with active chlorine species becomes faster in acid than in alkaline media because of the higher standard potentials of Cl₂ (E° = 1.36 V/SHE) and HClO (E° = 1.49 V/SHE) compared to that of ClO⁻ (E° = 0.89 V/SHE). However, results of Fig. 1 evidence a fast, quite similar destruction of paracetamol in both acid and alkaline real water matrices, probably due to the great quantities of active chlorine species produced at the high current density of 50 mA·cm⁻².

Note that the oxidation of Cl⁻ at the BDD anode can also yield other chlorine oxyanions like ClO₂⁻ from Reaction (8), ClO₃⁻ from Reaction (9) and/or ClO₄⁻ from Reaction (10) [7, 41-44]:

\[
\begin{align*}
\text{ClO}_2 + \text{H}_2\text{O} & \rightarrow \text{ClO}_2^- + 2\text{H}^+ + 2e^- \quad (8) \\
\text{ClO}_3^- + \text{H}_2\text{O} & \rightarrow \text{ClO}_3^- + 2\text{H}^+ + 2e^- \quad (9) \\
\text{ClO}_4^- + \text{H}_2\text{O} & \rightarrow \text{ClO}_4^- + 2\text{H}^+ + 2e^- \quad (10)
\end{align*}
\]

The existence of such reactions during the oxidation of Cl⁻ was confirmed from the ion chromatography analysis of the real water matrix with 1 mg·L⁻¹ paracetamol after 240 min of electro-oxidation in a BDD/Pt cell at 50 mA·cm⁻², which revealed about 90% decreases in the initial concentration of Cl⁻ along with the formation of near 20% ClO₂⁻.

The influence of current density on the decay of 1 mg·L⁻¹ paracetamol in synthetic and real aqueous matrices of pH 3.0 using the BDD/Pt cell is illustrat-
ed in Fig. 2. A very slow decay in concentration can be observed in ultrapure water at a low j of 10 mA · cm², only achieving 64% reduction after 60 min of electro-oxidation. Compared to Fig. 1, the removal rate of the pharmaceutical at 10 mA cm² is much lower than that at 50 mA · cm², where it disappears in 45 min, which can be related to the concomitant lower production of BDD(·OH) from Reaction (1). Nevertheless, it is well known²⁷-¹¹ that the increase in j causes a progressive decrease in the relative amount of BDD(·OH) that is available for the degradation of the organic molecules, which is due to the higher increase in rate of some waste reactions; for example, its anodic oxidation to O₂ from Reaction (11), its dimerization to H₂O₂ from Reaction (12), and its reaction with H₂O₂ generating the weaker oxidant hydroperoxyl radical (BDD(HO₂·)) from Reaction (13). The relative proportion of BDD(·OH) can also diminish by the larger enhancement of Reactions (14) and (15) at high j, giving rise to persulfate ion (SO₄²⁻) and oxygen, respectively.

\[
2\text{BDD}(\cdot\text{OH}) \rightarrow 2\text{BDD} + \text{O}_2 + 2\text{H}^+ + 2e \quad (11)
\]

\[
2\text{BDD}(\cdot\text{OH}) \rightarrow 2\text{BDD} + \text{H}_2\text{O}_2 \quad (12)
\]

\[
\text{BDD}(\cdot\text{OH}) + \text{H}_2\text{O}_2 \rightarrow \text{BDD}(\text{HO}_2^\cdot) + \text{H}_2\text{O} \quad (13)
\]

\[
2\text{SO}_4^{2-} \rightarrow \text{S}_2\text{O}_8^{2-} + 2e \quad (14)
\]

\[
3\text{H}_2\text{O} \rightarrow \text{O}_2 + 6\text{H}^+ + 6e \quad (15)
\]

As shown in Fig. 2, the abatement of paracetamol at 10 mA · cm² is much faster in the real water matrix, reaching its overall destruction in ca. 60 min, as a result of the combined action of active chlorine species (mainly Cl₂/HClO) and BDD(·OH). This suggests that the former species have a great contribution to the degradation of the organic molecule, due to: (i) their high oxidation power and (ii) the fact that they act in the bulk, instead of being confined in the anode vicinity, in contrast to BDD(·OH). Under these conditions, it should be highlighted that both oxidants can also attack the NOM contained in the real water matrix, as will be discussed below and, consequently, their oxidation ability is lower than using ultrapure water as solvent. Fig. 2 highlights that, at 50 mA · cm², the presence of 0.05 mol · L⁻¹ Na₂SO₄ in the real water matrix does not modify the rate of paracetamol decay, again yielding total removal in 5 min. This corroborates the predominant role of active chlorine species in the destruction of this pharmaceutical, since the decay kinetics is not affected by the plausible decrease in BDD(·OH) content arising from the acceleration of SO₄²⁻ formation from Reaction (14) in the presence of 0.05 mol · L⁻¹ Na₂SO₄. It is worth mentioning that the abatement of paracetamol concentration in all the above assays did not follow any kinetic equation related to simple order reactions, thus suggesting the existence of a complex oxidation route for this drug.

![Fig. 2 Time course of paracetamol concentration for the electro-oxidation of 100 mL of 1 mg · L⁻¹ of pharmaceutical in solutions of pH 3.0 and 25 °C using a BDD/Pt cell ((+) Ultrapure water with 0.05 mol · L⁻¹ Na₂SO₄, 10 mA · cm², (●) real water matrix, 10 mA · cm², (●) real water matrix, 50 mA · cm², and (●) real water matrix with 0.05 mol · L⁻¹ Na₂SO₄, 50 mA · cm²)](image)

### 2.2 Electro-Oxidation of Solutions Containing Salicylic Acid Alone

Aiming to confirm the above degradation behavior found for paracetamol, salicylic acid was treated under comparable conditions, so as to ascertain if it can be generalized for the electro-oxidation of low amounts of aromatic drugs with a BDD anode. As can be seen in Fig. 3, the decay of 1 mg · L⁻¹ salicylic acid in ultrapure water with 0.05 mol · L⁻¹ Na₂SO₄ of pH 3.0 using the BDD/Pt cell was very slow, being faster for 50 mA · cm² than for 10 mA · cm². This is expected from the greater production of the main oxido...
dant BDD(·OH) at higher $j$ values, as pointed out above. The fact that the total destruction of this compound takes place in 60 min at 50 mA cm$^{-2}$ (see Fig. 3), a time longer than 45 min required for 1 mg L$^{-1}$ paracetamol under the same conditions (see Fig. 1), is indicative of a slightly slower reaction kinetics between BDD(·OH) and salicylic acid. One can then infer that this radical is potent enough to completely remove low concentrations of aromatic pharmaceuticals in short electrolysis times operating at sufficiently high $j$ values.

When electro-oxidation was performed using the real water matrix, the decay in concentration of salicylic acid was strongly accelerated by its parallel oxidation with active chlorine species produced from Cl$^-$ oxidation by Reaction (5) and thereafter. The fact that the total destruction of this compound takes place in 60 min at 50 mA cm$^{-2}$ (see Fig. 3), a time longer than 45 min required for 1 mg L$^{-1}$ paracetamol under the same conditions (see Fig. 1), is indicative of a slightly slower reaction kinetics between BDD(·OH) and salicylic acid. One can then infer that this radical is potent enough to completely remove low concentrations of aromatic pharmaceuticals in short electrolysis times operating at sufficiently high $j$ values.

When electro-oxidation was performed using the real water matrix, the decay in concentration of salicylic acid was strongly accelerated by its parallel oxidation with active chlorine species produced from Cl$^-$ oxidation by Reaction (5) and thereafter. Fig. 3 highlights that by simply working at a low $j$ of 10 mA cm$^{-2}$, this compound was so rapidly destroyed that disappeared in only 7 min. Note that this time is much shorter than 60 min needed for the total abatement of paracetamol under the same conditions (see Fig. 2), evidencing the much faster reaction kinetics between generated active chlorine species (mainly Cl$_2$/HClO) and salicylic acid. Fig. 3 also shows that increasing $j$ from 10 to 50 mA cm$^{-2}$ in this medium only yielded a slight acceleration of its removal rate, disappearing in 5 min, in either the absence or the presence of 0.05 mol L$^{-1}$ Na$_2$SO$_4$. These findings allow concluding that in this system, salicylic acid is mainly attacked by active chlorine species, with small contribution of BDD(·OH), so that a $j$ value as low as 10 mA cm$^{-2}$ is enough to rapidly remove this drug from the medium. In all the experiments, its reaction with generated oxidants was quite complex since it did not follow any kinetic equation related to simple order reactions.

The aforementioned results demonstrate that the current density needed to destroy low contents of aromatic pharmaceuticals in a real water matrix by electro-oxidation with a BDD anode exhibits a great dependence on their reactivity with generated oxidants. Thus, the high oxidation ability of active chlorine species, as well as their distribution in the whole solution bulk that differs from confined BDD(·OH), may enhance the removal of these contaminants allowing the use of lower $j$ values than in non-chloride media to attain their overall disappearance in shorter electrolysis times.

![Fig. 3 Removal of salicylic acid with electrolysis time for the electro-oxidation with a BDD/Pt cell of 100 mL of 1 mg L$^{-1}$ of pharmaceutical in solutions of pH 3.0 and 25 °C: (+) Ultrapure water with 0.05 mol L$^{-1}$ Na$_2$SO$_4$, 10 mA cm$^{-2}$, (x) ultrapure water with 0.05 mol L$^{-1}$ Na$_2$SO$_4$, 50 mA cm$^{-2}$, (●) real water matrix, 10 mA cm$^{-2}$, (▼) real water matrix, 50 mA cm$^{-2}$, and (●) real water matrix with 0.05 mol L$^{-1}$ Na$_2$SO$_4$, 50 mA cm$^{-2}$](image)

### 2.3 Electro-Oxidation of Multicomponent Solutions with Paracetamol and Salicylic Acid

The electro-oxidation for mixtures of 1 mg L$^{-1}$ of both, paracetamol and salicylic acid, in synthetic and real matrices was studied to clarify the influence of generated oxidants once it is in competition with the other one. In view of the very fast abatement of salicylic acid in the real water matrix (see Fig. 3), only the changes in paracetamol decay in drug mixture solutions will be described below.

Fig. 4A shows a quicker destruction of paracetamol in a BDD/Pt cell in both synthetic and real solutions when $j$ rises from 10 to 50 mA cm$^{-2}$ as a result of the larger production of BDD(·OH) and/or active chlorine species due to the acceleration of Reactions (1) and (5), respectively. As expected from the above results, the drug was more rapidly removed in the real conditions.
water matrix because of the higher reactivity of active chlorine species formed in this medium. However, comparison in results of Fig. 4A with those of Figs. 1 and 2 for single paracetamol solutions allows inferring a slower decay of this compound in all the mixtures of both pharmaceuticals. For example, after 30 min of electrolysis at 10 mA · cm⁻² in the synthetic medium, its concentration was reduced by 48% in the single solution (Fig. 2) and by only 37% in the drug mixture (Fig. 4A), whereas at 50 mA · cm⁻² in the real water matrix, the time for its overall disappearance was prolonged from about 5 min in the single solution (Fig. 1) to near 8 min in the drug mixture (Fig. 4A). This phenomenon can be ascribed to the generation of similar amounts of the corresponding oxidants in each medium under comparable conditions, thereby being competitively consumed by both pharmaceuticals when mixed, so that their individual reaction rates with respect to that of the single drug solutions become slower.

The oxidation ability of other reactive oxygen species (ROS) like H₂O₂ and HO₂⁻ produced during electrolysis from Reactions (12) and (13), respectively, on paracetamol abatement in drug mixture solutions was assessed using a BDD/air-diffusion cell. Under these conditions, H₂O₂ is generated at the cathode from Reaction (2) and accumulated to a large extent in the medium, while it can be oxidized at the anode yielding HO₂⁻.¹⁸⁻²³

\[
\text{H}_2\text{O}_2 \rightarrow \text{HO}_2^- + \text{H}^+ + e \quad (16)
\]

The production of HO₂⁻ from Reaction (16) is much higher than from Reaction (13) due to the much greater generation of H₂O₂ at the cathode from Reaction (2) than at the anode from Reaction (12).²⁷

As an example, Fig. 4B depicts the decay in paracetamol concentration in ultrapure water with 0.05 mol · L⁻¹ Na₂SO₄ and in real water matrix using a BDD/air-diffusion cell at 10 mA · cm⁻². Comparison of these data with those obtained for the BDD/Pt cell under the same conditions (Fig. 4A) corroborates the existence of a slightly quicker destruction of the drug in the BDD/air-diffusion cell. Thus, at 60 min of electro-oxidation of the synthetic solution the paracetamol decay increased from 43% for the BDD/Pt cell (Fig. 4A) to 57% for the BDD/air-diffusion cell (Fig. 4B). The drug abatement was more enhanced in the real water matrix, although in both cells the time needed for its total removal was same (60 min as seen in Fig. 4A and B). These results evidence the low oxidation power of H₂O₂ and HO₂⁻ to destroy paracetamol, thereby confirming that BDD(·OH) is the main ROS in the electro-oxidation process.

Fig. 4 Change of paracetamol concentration with electrolysis time upon electro-oxidation of 100 mL pharmaceutical mixtures containing 1 mg · L⁻¹ of this drug and 1 mg · L⁻¹ salicylic acid at pH 3.0 and 25 °C (In plot A, BDD/Pt cell: Ultrapure water with 0.05 mol · L⁻¹ Na₂SO₄ at (+) 10 mA · cm⁻² and (•) 50 mA · cm⁻²; real water matrix at (□) 10 mA · cm⁻² and (△) 50 mA · cm⁻². In plot B, BDD/air-diffusion cell at 10 mA · cm⁻²: (+) Ultrapure water with 0.05 mol · L⁻¹ Na₂SO₄ and (■) real water matrix)

2.4 Electro–Fenton Treatment

The performance of the electro-Fenton process was assessed by adding 0.15 mmol · L⁻¹ Fe²⁺ to the solutions of pH 3.0 degraded in the BDD/air-diffusion
cell. The destruction of 1 mg·L⁻¹ paracetamol in the real water matrix up to overall disappearance after 60 min at 10 mA·cm⁻² is shown in Fig. 5. Surprisingly, the removal rate of this drug by electro-Fenton was only slightly higher than that found by electro-oxidation at 10 mA·cm⁻² (Fig. 2), in contrast to previous work[25-26] where the former method showed a much higher efficiency owing to the significantly faster reaction of paracetamol with homogeneous ·OH formed from Fenton’s reaction (3) than with BDD(·OH) in 0.05 mol·L⁻¹ Na₂SO₄. This phenomenon can be mainly explained by the inhibitory effect of Cl⁻ presented in the real water matrix on generated ·OH, giving rise to weak active chlorine species like ClO₂⁺, Cl₂⁺, Cl₃⁺, and Cl₃O via the following reactions[43-46]:

\[ \cdot \text{OH} + \text{Cl}^- \leftrightarrow \text{ClO}_2^+ \]  
\[ \text{ClO}_2^+ + \text{H}^+ \rightarrow \text{Cl}^+ + \text{H}_2\text{O} \]  
\[ \text{ClO}_2^+ + \text{Cl}^- \rightarrow \text{Cl}_2^+ + \text{OH}^- \]  
\[ \text{Cl}^- + \text{Cl}^- \leftrightarrow \text{Cl}_2^+ \]  
\[ 2\text{Cl}_2^+ \rightarrow \text{Cl}_3^0 + 2\text{Cl}^- \]  

The inhibition by Cl⁻ was confirmed from the similar paracetamol decay found for a drug mixture in synthetic and real solutions, as shown in Fig. 5. The enhancement of drug removal observed for its electro-oxidation in the real water matrix in Fig. 4 ascribed to the parallel action of active chlorine species was not produced in electro-Fenton because of the destruction of ·OH with Cl⁻. The presence of Cl⁻ in the real water matrix is then beneficial to increase the oxidation power of electro-oxidation, but clearly detrimental for electro-Fenton since it reduces its oxidation ability.

2.5 DOC Decay and Evolution of Oxidation By-Products in the Real Water Matrix

In natural and engineered systems, the reactivity between NOM and some oxidants such as ·OH is of great importance since these natural organic compounds may act as radical scavengers, therefore altering the performance of the AOPs[47]. The change of DOC with time for those trials shown above that were performed in real water matrix was determined to assess the degree in mineralization of its organic pollutants by the EAOPs tested. The initial DOC of 8.0 mg·L⁻¹ for the real water (Tab. 1) was increased to about 8.6 mg·L⁻¹ for the single drug solutions and to ca. 9.3 mg·L⁻¹ for the multicomponent solutions. The results obtained for selected experiments are depicted in Fig. 6. As can be seen, the use of 10 mA·cm⁻² in electro-oxidation yielded a very poor mineralization, reaching 15% of DOC decay at 180 min as a consequence of the low amounts of BDD(·OH) and active chlorine species produced at this low j. The mineralization rate of the polluted water was slightly enhanced by means of electro-Fenton, which only attained 23% DOC reduction after being electrolyzed for 180 min at 10 mA·cm⁻² and 51% DOC removal after the application of 50 mA·cm⁻² for 120 min (Fig. 6). The slightly better oxidation ability of electro-Fenton compared to electro-oxidation is due to the parallel destruction of organics with ·OH although its production from Fenton’s reaction (3) is partially inhibited by Cl⁻, as stated above. In contrast, Fig. 6 shows that at 50 mA·cm⁻² a high mineralization degree was already reached in electro-oxidation, being near 90% at 240 min of electrolysis, and even 95% at 180 min by the addition of 0.05 mol·L⁻¹ Na₂SO₄ probably due to the additional oxidation of pollutants by the larger quantities of S₄O₆²⁻ formed from Reaction (14). High j
values and high contents of Cl\textsuperscript{-} and SO\textsubscript{4}\textsuperscript{2-} are then key factors to obtain an almost total mineralization of the dissolved NOM in the real water matrix upon electro-oxidation.

Reversed-phase HPLC analysis of single paracetamol solutions treated by electro-oxidation revealed the formation of traces of \textit{p}-benzoquinone (<0.2 mg·L\textsuperscript{-1}), which persisted for long time, for example, up to 240 min at 50 mA·cm\textsuperscript{-2}. This derivative can arise from the oxidation of hydroquinone, which is formed from the paracetamol hydroxylation with cleavage of the C(1)—N bond\textsuperscript{[11]}. On the other hand, ion chromatograms of all treated paracetamol solutions exhibited traces of NO\textsubscript{3}\textsuperscript{-} and NH\textsubscript{4}\textsuperscript{+} ions, presumably produced from the drug degradation\textsuperscript{[11,26]} because the TN measured for the trials with the real water matrix did not vary with electrolysis time. This indicates that N-compounds contained in the real water matrix are very recalcitrant since they are destroyed by neither hydroxyl radicals nor active chlorine species generated in electro-oxidation and electro-Fenton.

3 Conclusions

It has been demonstrated that 1 mg·L\textsuperscript{-1} paracetamol and/or 1 mg·L\textsuperscript{-1} salicylic acid contained in a real water matrix at pH 3.0 are destroyed by electro-oxidation with a BDD anode. Their decay is faster in this medium than in a synthetic solution with 0.05 mol·L\textsuperscript{-1} Na\textsubscript{2}SO\textsubscript{4} due to their favored reaction in the bulk with highly oxidizing active chlorine species produced from the oxidation of Cl\textsuperscript{-} ion. The increase in \textit{j} from 10 to 50 mA·cm\textsuperscript{-2} causes a large enhancement in the removal of both compounds, which disappear in about 5 min, as a result of the high production of oxidants. This behavior also occurs using the real water mixture at natural alkaline pH. When the drug mixture is treated, the competitive attack of generated oxidants over them decelerates their abatement. Comparative assays with BDD/Pt and BDD/air-diffusion cells have shown a very low oxidation ability of other ROS like H\textsubscript{2}O\textsubscript{2} and HO\textsubscript{2}● in comparison to BDD (•OH) in electro-oxidation. Furthermore, only a slight acceleration of drug removal has been found in the real water matrix using electro-Fenton with a BDD anode, owing to the partial destruction of homogeneous •OH formed from Fenton’s reaction (3) with Cl\textsuperscript{-}. High mineralization degrees can be reached at a high \textit{j} of 50 mA·cm\textsuperscript{-2}, attaining an almost total mineralization with 95% DOC (mainly coming from NOM) reduction for the real water matrix with 1 mg·L\textsuperscript{-1} paracetamol and 0.05 mol·L\textsuperscript{-1} Na\textsubscript{2}SO\textsubscript{4} after 180 min of electro-oxidation. Traces of \textit{p}-benzoquinone as well as NO\textsubscript{3}\textsuperscript{-} and NH\textsubscript{4}\textsuperscript{+} ions have been detected during the electro-oxidation of single paracetamol solutions. However, the TN of the real water matrix is not reduced in none of the trials, which suggests that its N-compounds remain stable upon treatment by these EAOPs under the present conditions.

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![Fig. 6 Normalized DOC decay vs. electrolysis time for 100 mL real water matrix at pH 3.0 and 25 °C ( □ , △ , ●) 1 mg·L\textsuperscript{-1} paracetamol, by electro-oxidation; (○) 1 mg·L\textsuperscript{-1} paracetamol + 1 mg·L\textsuperscript{-1} salicylic acid, by electro-Fenton. Applied current density: (□ , ○) 10 mA·cm\textsuperscript{-2} and (△ , ●) 50 mA·cm\textsuperscript{-2}. (●) In the presence of 0.05 mol·L\textsuperscript{-1} Na\textsubscript{2}SO\textsubscript{4}.


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掺硼金刚石阳极电氧化和电芬顿工艺处理真实水体中低含量药物的研究

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Abstract: This study reports the use of boron-doped diamond (BDD) electrodes for the electrochemical treatment of model solutions containing low levels of pharmaceuticals in real water. Results from experiments conducted at constant current density in BDD/Pt electrodes for chronoamperometric experiments and in BDD/air-diffusion electrodes for Fenton’s process show that the oxidation of acetaminophen and salicylic acid proceeded more effectively in real water compared to ultrapure water containing 0.05 mol L⁻¹ Na₂SO₄. This is attributed to the formation of reactive halogen species, which accelerated the degradation of the pollutants. When both drugs were treated in real water, the competing effects of the two oxidants slowed down the degradation of the pollutants, whereas Fenton’s process accelerated the degradation of the drugs in real water due to the formation of hydroxyl radicals. Upon increasing the current density, real water showed similar results to ultrapure water. No significant differences in the degradation of the mixtures were observed.

Keywords:掺硼金刚石阳极; 电氧化; 电芬顿; 药物; 真实水体; 水杨酸