



UVA-UVB activation of hydrogen peroxide and persulfate for advanced oxidation processes: Efficiency, mechanism and effect of various water constituents

Wenyu Huang, Angelica Bianco, Marcello Brigante, Gilles Mailhot

► To cite this version:

Wenyu Huang, Angelica Bianco, Marcello Brigante, Gilles Mailhot. UVA-UVB activation of hydrogen peroxide and persulfate for advanced oxidation processes: Efficiency, mechanism and effect of various water constituents. *Journal of Hazardous Materials*, 2018, 347, pp.279 - 287. 10.1016/j.jhazmat.2018.01.006 . hal-01772765

HAL Id: hal-01772765

<https://hal.science/hal-01772765>

Submitted on 17 Dec 2020

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

UVA-UVB activation of hydrogen peroxide and persulfate for Advanced Oxidation Processes: Efficiency, mechanism and effect of various water constituents

Wenyu Huang^{a,b*}, Angelica Bianco^a, Marcello Brigante^{a*}, Gilles Mailhot^a

^a Université Clermont Auvergne, CNRS, SIGMA Clermont, Institut de Chimie de Clermont-Ferrand,
F-63000 Clermont–Ferrand, France.

^b School of the Environment, Guangxi University, Nanning 530004, China

*Corresponding authors : huangwenyu@gxu.edu.cn (WH) and marcello.brigante@uca.fr (MB)

Abstract

In the present work we investigate the activation efficiency of H_2O_2 and $\text{S}_2\text{O}_8^{2-}$ using UVA and UVB radiation. Bisphenol-A (BPA) is used as model pollutants to estimate the oxidative process efficiency in simulated and real sewage treatment plant waters. Particular attention is paid to the BPA removal efficiency and to the radical mechanism involvement considering the effect of typical

inorganic water constituents (carbonates and chloride ions) and organic matter. Despite a detrimental effect observed when carbonate ions are in solution using both hydrogen peroxide and persulfate, the presence of high chloride ions concentration was found to improve BPA removal using $S_2O_8^{2-}$ as radical precursor. This enhancement, investigated combining chemical kinetic model approach and laser flash photolysis experiments, is attributed to the formation of hydroxyl radical and chlorine radical species from sulfate radical. Different transformation products are identified by means of GC-MS and HPLC-MS analyses. Moreover, experiments using sewage treatment plant water (STPW) spiked with BPA are performed in order to assess the efficiency of oxidative processes in a simulated treatment systems activated using UVA+UVB radiations.

Keywords: AOPs, pollutant degradation, chloride ions effect, hydroxyl and sulfate radical, Bisphenol A

1. Introduction

Treatment efficiency of recalcitrant organic pollutants in aqueous effluents using oxidant species precursors such as hydrogen peroxide (H_2O_2) and persulfate ($S_2O_8^{2-}$) has been investigated under UVC radiation [1-5]. UVC lamps are mercury made, fragile and they require considerable more energy than UVA, UVB lamps [6] which can also be substituted by LED to reduce the operating cost. UVC-LED efficiency is less than 3 % making their utilization not suitable for long time wastewater treatments. For these reasons, application of UVA and UVB radiation to improve water

depollution processes can offer many advantages. Following this idea, iron-complexes are recently used to initiate or improve pollutant removal using H_2O_2 and $\text{S}_2\text{O}_8^{2-}$ activation in the presence of solar spectrum UV radiation (UVA and UVB) [7-11]. In this photo-activated processes, generated hydroxyl and sulfate radicals (HO^\bullet and $\text{SO}_4^{\bullet-}$) can react with a wide range of organic pollutants due to their high redox potential. And the redox potentials vs NHE and in standard conditions of HO^\bullet and $\text{SO}_4^{\bullet-}$ is 2.7V and 2.6V respectively [12]. Therefore, application H_2O_2 or $\text{S}_2\text{O}_8^{2-}$ photoactivation in wastewater treatment is now taken into account for emerging contaminant abatement. Comparing to HO^\bullet based processes, $\text{SO}_4^{\bullet-}$ based methods from activation Persulfate (PS, $\text{S}_2\text{O}_8^{2-}$) have a series of advantages, and attract many researches [13]. However, the presence of organic and inorganic species (such as carbonate and chloride ions) in natural and wastewater may strongly influence the efficiency of oxidation treatment [14, 15]. For this reason, several investigations providing a better understanding of natural occurring ions effect on the oxidative process are reported [16-18]. Generally, chloride ions (Cl^-) can significantly inhibit efficiency of UVC based activation of H_2O_2 and $\text{S}_2\text{O}_8^{2-}$ due to their radical scavenging effect [19, 20]. However, in the presence of chloride ions, generated reactive species such as chlorine radical (Cl^\bullet) and dichlorine radical anion ($\text{Cl}_2^{\bullet-}$) could also react with organic pollutants being more selective [16]. On the other hand, it was suggested that the chain reaction triggered by Cl^- and $\text{SO}_4^{\bullet-}$ can result in the production of HO^\bullet [21, 22] and conflicting results are reported in different studies on this topic. For example, different researchers [17, 22, 23] reported that the presence of Cl^- would inhibit the oxidation of organics, while in other investigations, a negligible effect of chloride ions on the

UVC-H₂O₂ treatment performances was observed [24]. Moreover, using SO₄^{•-} based oxidation, pollutant's degradation mechanism was found to be modified due to the presence of chloride ions [25, 26].

Inorganic carbon (HCO₃⁻/CO₃²⁻, pKa =10.33) is also an important water constituent that considerably influence the oxidation process. In fact, reactivity of HCO₃⁻/CO₃²⁻ with HO[•] or SO₄^{•-} generates carbonate radical anion (CO₃^{•-}) [27, 28] which is generally less reactive toward organic molecules (the second order rate constant with organic compounds in the range ~ 10⁵ – 10⁹ M⁻¹ s⁻¹) than HO[•] or SO₄^{•-}. However, CO₃^{•-} can efficiently react with electron-rich atoms, such as N and S-containing compounds, phenols and inorganic compounds [15, 29].

Besides, even if the effect of single water constituent could mutually offset or superimpose, the contribution on the oxidation process of different water matrix as a whole was also investigated in several previous researches [30-34]. Kattel et coworkers found strong effect of the natural water composition on the oxidative degradation of emerging micropollutant acesulfame using UVA-induced H₂O₂/Fe²⁺ and S₂O₈²⁻/Fe²⁺ processes [35]. Due to the complexity of different water matrix, the efficiency of HO[•] or SO₄^{•-} based oxidation processes (also called AOPs: advanced oxidation processes) strongly depends on the investigated medium and on the target pollutant. In the present work, UVA + UVB (UVA/B) radiation is used to activate H₂O₂ and S₂O₈²⁻ using Bisphenol A (BPA) as target compound. BPA degradation efficiency is investigated in the presence of natural and wastewaters occurring inorganic constituents such as chloride and hydrogenocarbonate/carbonate ions and in real sewage treatment plant water (STPW). Bimolecular rate constants of reaction of

BPA and STPW constituents with radicals such as HO^\bullet , $\text{SO}_4^{\bullet-}$, $\text{Cl}_2^{\bullet-}$ and $\text{CO}_3^{\bullet-}$ are estimated through laser flash photolysis experiments and used in a chemical kinetic model. Finally, radical species involvement in the presence of different water constituents and impact on the oxidative process are discussed.

2. Materials and methods

2.1 Chemicals

Hydrogen peroxide (H_2O_2) (30% in water) is purchased from Fluka, France. Bisphenol A (BPA), sodium persulfate ($\text{Na}_2\text{S}_2\text{O}_8$), sodium chloride (NaCl) and sodium hydrogen carbonate (NaHCO_3) are obtained from Sigma, France. Perchloric acid (HClO_4) and sodium hydroxide (NaOH) are used to adjust the pH of the solutions. 1 mM buffer solution with sodium dihydrogen phosphate (NaH_2PO_4) and sodium hydrogen phosphate (Na_2HPO_4) is used to keep pH value constant at 8.0 during the irradiation time without otherwise stated. All chemicals are used without further purification.

2.2 Irradiation experiments and analysis

Irradiations are performed in a cylindrical Pyrex reactor (500 mL) placed in a rectangular box, equipped, on the top, with four fluorescence lamps whose emitting wavelength is larger than 275 nm (UVA/B radiation). The polychromatic emission spectrum of the irradiation system reaching the solution is recorded using an optical fiber with a charge coupled device (CCD) spectrophotometer

(Ocean Optics USD 2000 + UV-vis) which is calibrated using a DH-2000-CAL Deuterium Tungsten Halogen reference lamp. The total irradiance between 275 and 400 nm reaching the solution is then estimated to be $976 \mu\text{W cm}^{-2}$. In Figure SM1 emission spectrum of lamps and overlap with absorption spectra of H_2O_2 and $\text{S}_2\text{O}_8^{2-}$ are presented. All experiments are performed at room temperature ($293 \pm 2 \text{ K}$) using a circulation cooling system.

BPA solution ($43.8 \mu\text{M}$ corresponding to 10 mg L^{-1}) is magnetically stirred with a magnetic bar during the reaction to ensure the homogeneity of solution. All irradiation experiments are performed using a phosphate buffer (total buffer concentration of 0.1 mM) without otherwise stated. At fixed interval times, 1 mL of sample is withdrawn and analyzed by ultra performance liquid chromatography (UPLC, ACQUITY, Waters, USA) equipped with photodiode array detector (PDA). The flow rate is 0.25 mL min^{-1} and the mobile phase is a mixture of water and methanol (40/60, v/v). The column is a Zorbax C18 of $150 \times 4.6 \text{ mm}$ with particle size $5 \mu\text{m}$. BPA disappearance is followed at 225 nm .

HPLC-MS and GC-MS conditions used for degradation identification are reported in the supplementary material section.

The pseudo-first-order decay of BPA is determined by plotting BPA disappearance vs time and using following relation:

$$\frac{[\text{BPA}]_t}{[\text{BPA}]_0} = \exp(-k_{app}t)$$

Where $[\text{BPA}]_0$ and $[\text{BPA}]_t$ are the initial and remaining concentrations of BPA at time t , k_{app} is the

pseudo-first-order apparent rate constant (s^{-1}). The initial transformation rate of BPA is R_{BPA} ($M s^{-1}$) $= k_{app} \times [BPA]_0$. Degradation efficiency (DE) of considered system is determined using the following equation:

$$DE(\%) = \left[\left(\frac{[BPA]}{[BPA]_0} \right)_R / \left(\frac{[BPA]}{[BPA]_0} \right)_S - 1 \right] \times 100 \%$$

where $\left(\frac{[BPA]}{[BPA]_0} \right)_R$ and $\left(\frac{[BPA]}{[BPA]_0} \right)_S$ are the value of remaining BPA in solution in absence (R) and in the presence (S) of inorganic ion determined experimentally.

Sewage Treatment Plant Water (STPW) is obtained from urban sewage treatment plant of Clermont-Ferrand, France (April 29th, 2016) and main physico-chemical characteristics are reported in Table 1. STPW is used for photochemical experiments immediately after PTFE (0.45 μm size pore) filtration in order to avoid bacteriological and chemical modifications. All experiments are performed in triplicates in order to ensure the reliability of the results.

2.3 Laser flash photolysis

The laser flash photolysis apparatus has been previously described [36] and a brief description of radical species detection is given in this section. The fourth harmonic ($\lambda_{exc} = 266$ nm) of a Nd:YAG laser is used for the excitation (the energy of pulse is 40 mJ). An appropriate volume of chemical stock solutions (BPA, $S_2O_8^{2-}$, H_2O_2 , HCO_3^-/CO_3^{2-} , Cl^- and STPW) is mixed just before each experiment to obtain the desired mixtures and concentrations. All experiments are performed at ambient temperature (293 ± 2 K) and in aerated solutions. Hydroxyl radical (HO^\bullet) reactivity is

determined by using chemical competition kinetics with thiocyanate anion (SCN^-) and di-thiocyanate radical anion ($\text{SCN}_2^{\bullet-}$) specie, and the detection wavelength of $\text{SCN}_2^{\bullet-}$ was at 470 nm. Sulfate radical anion ($\text{SO}_4^{\bullet-}$) decay is followed at 450 nm corresponding to the maximum absorption of this species [37]. Dichlorine radical anion ($\text{Cl}_2^{\bullet-}$) and carbonate radical anion ($\text{CO}_3^{\bullet-}$) species are generated using electron transfer reaction between photogenerated sulfate radical and chloride/carbonate ions. The same approach has been previously used to generate $\text{Cl}_2^{\bullet-}$ in solution and follow the reactivity toward organic molecules [38]. Details concerning the second order rate constant determinations are reported in Figure SM2. The second order rate constant between selected radicals ($\text{SO}_4^{\bullet-}$, $\text{Cl}_2^{\bullet-}$ and $\text{CO}_3^{\bullet-}$) and BPA or STPW is determined from the regression lines of the logarithmic decays of radical transient (monitored at 450 nm, 340 nm and 600 nm respectively) as a function of the quencher concentration (for BPA) or normalised for the carbon concentration (for STPW). Each value is the average of 4 consecutive laser pulses and the reported error is $\pm 3\sigma$, obtained from the scattering of the experimental data from the fitting line.

2.4 Kinetic modelling

A kinetic modelling approach is used to estimate the radical species formation during irradiation time in the presence of chloride ions. Table SM1 shows the chemical reactions considered for kinetic modelling (reactions are reported from M1 to M21). The pseudo-first order decay and second-order rate constants are obtained from the literature or experimentally (see results and discussion section). The numerical differential equations of the reaction rates are integrated using

the ode15s solver function of Matlab. The initial species concentration is implemented in the m-file as the input data. Pseudo-first order constant for the formation of hydroxyl and sulfate radicals (k_1 and k_2) are determined under adopted photochemical conditions (irradiance and initial precursor concentrations of 2 mM).

3. Results and discussion

3.1 Comparison of UVA/B activation of H_2O_2 and $\text{S}_2\text{O}_8^{2-}$

In Figure 1 the disappearance of 43.8 μM BPA (expressed as $[\text{BPA}]/[\text{BPA}]_0$) is followed when 2 mM of H_2O_2 and $\text{S}_2\text{O}_8^{2-}$ are irradiated using UVA/B using phosphate buffered and unbuffered solutions at pH 8.0 (which is representative of STPW pH value). Despite no direct photolysis of BPA is observed in pure water (milli-Q), in the presence of persulfate, the degradation rate results to be faster than in the presence of hydrogen peroxide. Such degradation was not observed under dark condition (direct oxidation) indicating that hydrogen peroxide and persulfate can be photochemically activated leading to the degradation of BPA through formation of highly oxidative radical species. In UVA/B– H_2O_2 system, the faster BPA degradation observed at pH 8.0 without buffer can be explained considering the change of pH value during irradiation. In fact, the solution pH slowly decreases up to pH 6.4 at the end of the experiment. This modification is correlated to the radical efficiency oxidation increase as reported in Figure SM3 (BPA degradation rate (R_{BPA}) increases from $\sim 1.2 \times 10^{-9} \text{ M s}^{-1}$ at pH 10 to $\sim 5.3 \times 10^{-9} \text{ M s}^{-1}$ at pH 4.6). In fact, it has been reported that the standard redox potential of HO^\bullet in acidic pH is about 2.4-2.7 eV which is higher

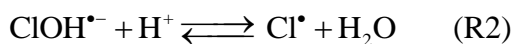
than that in alkaline pH (1.9–2.0 eV) [39]. On the contrary, in the UVA/B–S₂O₈²⁻ system, where sulfate radical are generated, a similar value for the BPA degradation rate is found between 4.6 and 9.0 ($\sim 8 \times 10^{-9} \text{ M s}^{-1}$). Such effect can be explained considering that persulfate radical redox potential does not depend on the pH [40]. Concerning the possible effect of phosphate buffer constituents, the bimolecular rate constants between photogenerated hydroxyl or sulfate radicals and HPO₄²⁻/H₂PO₄⁻ are $\leq 10^6 \text{ M}^{-1} \text{ s}^{-1}$ [41] so we can argue that the quenching effect of buffer could be negligible under adopted experimental conditions.

3.2 Effect of chloride ions

3.2.1 UVA/B–H₂O₂ system

In Figure 2, the effect of chloride ions on the BPA degradation efficiency using 2 mM of H₂O₂ is investigated at pH 4.6 and 8.0 under irradiation. At pH 8.0, slight inhibition of chloride ions (from 2 to 10 mM) is observed compared to the degradation using hydrogen peroxide alone (increasing the chloride ion concentration up to 100 mM the inhibition reaches ~15%). However, the detrimental effect of chloride ions is more marked at pH 4.6 where the BPA removal efficiency strongly decreases in the presence of 5 mM Cl⁻. The different BPA degradation trend at pH 8.0 and 4.6 can be justified considering that photogenerated hydroxyl radicals are in equilibrium with ClOH^{•-} specie in the presence of chloride ions (R1 and see reaction M13 in table SM1 for reactivity constant value) leading to the formation of Cl[•] only in acidic media (R2).





On the other hand, at pH 8.0 only a negligible amount of hydroxyl radical are expected to be finally converted into Cl^{\bullet} compared to pH 4.6.

The effect of chloride ions on the radical species formation can be partially predicted using the chemical kinetic modelling approach that estimates a negligible effect of Cl^- on the hydroxyl radical formation rate at pH 8.0. Despite no significant inhibition of chloride ions on the hydroxyl radical concentration (the initial concentration of HO^{\bullet} decreases from $\sim 4.1 \times 10^{-13} \text{ M}$ to $\sim 3.9 \times 10^{-13} \text{ M}$ in the presence of 5 mM of Cl^- at pH 4.6 as reported in Fig. SM4), the amount of chlorine radical species such as Cl^{\bullet} , $\text{Cl}_2^{\bullet-}$ and $\text{ClOH}^{\bullet-}$ is strongly enhanced (Fig. SM5). In fact, the kinetic model predicts that $\text{Cl}_2^{\bullet-}$, which is about 2 times less reactive toward BPA than HO^{\bullet} , is the main specie present in solution (see Table 2) explaining the inhibition of BPA degradation when 5 mM of chloride ions are mixed to H_2O_2 .

3.2.2 UVA/B- $\text{S}_2\text{O}_8^{2-}$ system

The BPA degradation using UVA/B- $\text{S}_2\text{O}_8^{2-}$ system is found to be strongly influenced by chloride ions (Fig. SM6). Two contrasting effects can be observed increasing the chloride ion concentration: the first (from 0 to 2 mM of Cl^-) is the drastic decrease (up to 80%) of the degradation efficiency (DE), whereas the second ($[\text{Cl}^-] > 2 \text{ mM}$) is a positive effect up to the enhancement of BPA degradation observed in the presence of 100 and 200 mM of Cl^- (Fig. 3). This opposite trend (inhibition and enhancement) is investigated considering the radical species generation as a function

of chloride ions concentration and their reactivity with photogenerated sulfate radical. Without chloride, sulfate radicals are able to oxidize water and hydroxide ions leading to the formation of hydroxyl radical in solution (see reactions M3 and M4 in Table SM1). Considering the second order rate constant of these two reactions, we can argue that less than 0.4 % of sulfate radical are converted into hydroxyl radical (mainly through reaction M3). The kinetic model, considering the photolysis rate of persulfate obtained experimentally (Fig. SM7 and SM8), estimates a steady state concentration of sulfate and hydroxyl radicals ($[\text{SO}_4^{\bullet-}]_{\text{ss}}$ and $[\text{HO}^{\bullet}]_{\text{ss}}$) of 1.5×10^{-10} and 4.6×10^{-12} M when 2 mM of persulfate are irradiated using UVA/B radiation at pH 8.0 (Fig. 4). The presence of 2 mM of chloride ions is able to scavenge about 66% of sulfate radicals leading to the formation of Cl^{\bullet} which is a source of other radical species such as $\text{Cl}_2^{\bullet-}$ and $\text{ClOH}^{\bullet-}$ through a more complex reaction system (M13 to M15). Increasing the chloride ions concentration, despite no effect observed on the hydroxyl radical steady-state concentration, sulfate radical concentration decreases of about 2 order of magnitude (from 10^{-12} to 10^{-14} M). This trend can be explained considering that sulfate radicals are able to oxidize chloride ions into chlorine radicals (M16) that can be responsible for the BPA degradation increase observed at high chloride ions concentrations. For instance, the reactivity of Cl^{\bullet} with phenol has been estimated to be $2.5 \times 10^{10} \text{ M s}^{-1}$ [42]. BPA degradation enhancement observed when persulfate are irradiated in the presence of chloride ions can be also explained considering the formation of new reactive species (that are not considered by our kinetic model) such as hypochlorous acid/hypochlorite ions (HOCl/OCl^- $\text{pK}_a = 7.5$) as suggested by Yuan and co-workers [43]. Chlorine active species are responsible for direct oxidation of organic

molecules or represent a photochemical source of hydroxyl and chlorine radical under UV radiation [44, 45].

3.3 Effect of $\text{HCO}_3^-/\text{CO}_3^{2-}$

The effect of carbonates ($\text{HCO}_3^-/\text{CO}_3^{2-}$ pka = 10.33) is investigated for UVA/B activation of H_2O_2 and $\text{S}_2\text{O}_8^{2-}$ at pH 8.0 and pH 9.0 using buffer solutions (see Figure SM9). The presence of 5 mM of carbonates at pH 8.0 in UV- H_2O_2 system inhibits about 50 % of the BPA degradation (the pseudo first order decay decreases from $5.3 \times 10^{-5} \text{ s}^{-1}$ to $2.6 \times 10^{-5} \text{ s}^{-1}$) while the effect on the UV- $\text{S}_2\text{O}_8^{2-}$ system is estimate to be around 18 % (the pseudo first order decay decreases from $1.9 \times 10^{-4} \text{ s}^{-1}$ to $1.5 \times 10^{-4} \text{ s}^{-1}$).

The scavenging effect of carbonates on photo-activation of hydrogen peroxide and persulfate can be predicted considering the reactivity of hydroxyl and sulfate radicals toward $\text{HCO}_3^-/\text{CO}_3^{2-}$ ($k_{\text{HO}^\bullet, \text{HCO}_3^-} = 8.6 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$, $k_{\text{SO}_4^{\bullet-}, \text{HCO}_3^-} = 2.8 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$, $k_{\text{HO}^\bullet, \text{CO}_3^{2-}} = 3.5 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$, $k_{\text{SO}_4^{\bullet-}, \text{CO}_3^{2-}} = 6.1 \times 10^6 \text{ M}^{-1} \text{ s}^{-1}$) [21, 46] and BPA determined by laser flash photolysis experiments (see Table 2). In Figure 5 the inhibition of the reactivity toward BPA due to the presence of $\text{HCO}_3^-/\text{CO}_3^{2-}$

is estimated as the ratio between the pseudo-first order decays of considered radicals with

carbonates and BPA ($\frac{k'_{\text{Carbonates}}}{k'_{\text{BPA}} + k'_{\text{Carbonates}}} \times 100$) as a function of carbonates concentration and pH. It is

interesting to observe that, while for sulfate radical the inhibition is not affected by the pH of the solution, for hydroxyl radical a negative effect is correlated to the pH increase. In fact, after pH ~8.2

we expect the presence of CO_3^{2-} species, which have higher reactivity toward hydroxyl radical (at pH 9.5 ~13 % of CO_3^{2-} and 87% of HCO_3^- are present in solution).

However, generated carbonate radicals are expected to react with BPA partially counterbalancing the sink of hydroxyl and sulfate radicals. Interestingly, degradation profiles of BPA at pH 9.0 compared to the pH 8.0 (see Figure SM9) show a less pronounced negative effect of carbonates on the inhibition of BPA degradation. Such effect can be explained considering the reactivity between $\text{CO}_3^{\bullet-}$ and phenol/phenolate groups. Moore and co-workers reported that reactivity of $\text{CO}_3^{\bullet-}$ with phenolate is about 6 times higher than the value measured for phenol ($k_{\text{CO}_3^{\bullet-}, \text{Phenol}} = 3.9 \times 10^8 \text{ M}^{-1} \text{ s}^{-1}$ and $k_{\text{CO}_3^{\bullet-}, \text{Phenolate}} = 2.4 \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$) [47]. We can reasonably expect a similar trend for BPA where increasing formation of carbonate radical at higher pH can enhance degradation of target pollutant in water.

3.4 Transformation products identification

Different experiments are conducted using BPA concentration of 43.8 μM in the presence of *i*) hydrogen peroxide (2 mM), *ii*) hydrogen peroxide (2 mM) + chloride ions (100 mM), *iii*) persulfate (2 mM) and *iv*) persulfate (2 mM) + chloride ions (100 mM). Table SM2 summarizes the chemical structures of transformation products identified using HPLC-MS (negative mode) and GC-MS on the basis of molecular peak, fragmentation and literature data.

Interestingly, hydroxylated BPA products are mainly found by HPLC-MS indicating that first reaction step of sulfate and hydroxyl radicals is the hydrogen abstraction on the aromatic ring and

consequent water addition. Sharma et al. investigated the intermediates of BPA degradation using UVC-peroxymonosulfate system and considered that hydrogen abstraction is not the only process occurring, but one of the most important initial steps of BPA degradation [48]. The authors concluded that BPA is oxidized by $\text{SO}_4^{\bullet-}$ through electron transfer and then H abstraction generates a hydroxyl BPA radical. Similar transformation products are found in all conditions with aromatic ring scission derivatives, which are also normally found in BPA degradation pathway induced by HO^{\bullet} or $\text{SO}_4^{\bullet-}$ attack [49-51]. However, a chlorinated BPA product is detected only using UVA/B- $\text{S}_2\text{O}_8^{2-}$ in the presence of chloride ions. Chlorinated derivatives have been previously detected by Zafra et al using chlorination experiments [52] corroborating the hypothesis that reactivity of sulfate radical can lead, in the presence of chloride ions, to the formation of new species such as HOCl/OCl^- that are expected to directly oxidize BPA in solution. The formation of chlorinated derivative reasonably suggests that the presence of high Cl^- concentration would change the pathway of BPA degradation in the presence of sulfate radical, while this phenomenon is not observed in UVA/B- H_2O_2 system. Moreover, high mass values are found in HPLC-MS ($m/z = 453$) in the presence of sulfate radical anions suggesting the formation of dimeric molecules produced *via* radical-radical recombination.

3.5 BPA degradation in STPW

To test the feasibility and performances of UVA/B activation of hydrogen peroxide and persulfate in simulated treatment systems, STPW are spiked with 4.3 μM and 430 nM of BPA (1 and 0.1 mg L^{-1}

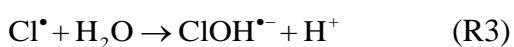
respectively) before irradiation in the presence of 2 mM of radical precursors. Due to the complexity of the STPW matrix, the inhibition effect of naturally constituents is investigated through several experiments. First, the degradation of BPA is carried out in STPW previously acidified using HClO_4 to $\text{pH} < 3$ in order to remove inorganic carbon (i.e. carbonate ions), and then basified back to pH around 8 (indicated above as STPW*). The purpose of this experiment is to discriminate the effect of organic and inorganic carbon on the degradation efficiency of considered system. These experiments show that the degradation of BPA in STPW and STPW* is almost the same; only a slight BPA inhibition (less than 5%) is observed in the presence of persulfate indicating a negligible effect of inorganic carbon (i.e. carbonates) on the degradation treatment efficiency (Fig. 6). Since the concentration of inorganic carbon is 4.16 mM in our STPW, it could be concluded that in this concentration level the degradation of organic compound is almost not affected by inorganic carbon. This trend can be also investigated considering the reactivity constant determined between hydroxyl and sulfate radicals and STPW normalizing the value to the carbon concentration (organic and inorganic). In Table 2 it is interesting to observe that reactivity constants using not modified STPW are about two times for hydroxyl radical and three times for sulfate radical lower than those determined after inorganic carbon removal (i.e. STPW*).

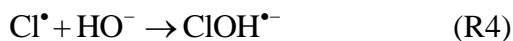
Considering the second order rate constant between hydroxyl radical and sulfate radical anion with STPW matrix, HCO_3^- (the main species present at pH 8.0) and BPA reported in this study (see Table 2), it is possible to investigate the formation pathways of photogenerated radicals in solution. For this purpose we considered that hydroxyl radicals (or sulfate radicals) can react with STPW

constituents or BPA in solution.

In the system UVA/B–H₂O₂, we can argue that about 98.4 % of hydroxyl radicals react with constituents of STPW (16.6 % of which accounted for by carbonates) while 1.6 % directly with BPA (Table 3). In the system UVA/B–S₂O₈²⁻ about 99.8 % of the sulfate radical reactivity is attributed to the STPW constituents, while only 0.2% is accounted by for BPA (Table 3). However, the BPA degradation results to be faster in the system UVA/B–S₂O₈²⁻ (Fig. 6).

The high efficiency degradation using UVA/B–S₂O₈²⁻ system could be explained considering that about 72 % of sulfate radicals react with chloride ions leading to the formation of chlorine radical which is a secondary source of hydroxyl radical and other chlorine reactive species in alkaline media. In order to assess the occurring reactive involvement using UVA/B–S₂O₈²⁻ process, a new experiment is performed spiking the solution with specific radical scavengers. STPW is spiked with 4.3 μM of BPA in the presence of 2-propanol (2-Pr) which is able to scavenge hydroxyl radical and sulfate radical ($k_{HO^{\bullet}, 2-Pr} = 1.9 \times 10^9 M^{-1} s^{-1}$, $k_{SO_4^{\bullet-}, 2-Pr} = 8.2 \times 10^7 M^{-1} s^{-1}$) and *tert*-butanol (*t*-But) which is more selective scavenger of hydroxyl radical ($k_{HO^{\bullet}, t-But} = 3.1 \times 10^9 M^{-1} s^{-1}$, $k_{SO_4^{\bullet-}, t-But} = 8.4 \times 10^5 M^{-1} s^{-1}$) [46, 53]. In the presence of 50 mM of *t*-But the degradation rate is inhibited of ~ 50 %, while, using 2-Pr, only 5% of BPA degradation is observed (Fig. 7). This result confirms that a significant fraction of sulfate radicals are converted into hydroxyl radical in STPW through a complex mechanism. In fact, chlorine radical can be converted into ClOH^{•-} (R3 and R4) which leads to the formation of hydroxyl radical in solution at pH 8.0 (see reaction R1).





Moreover, our conclusion are confirmed considering the high scavenging efficiency of 2-Pr and *t*-But on chlorine radical ($k_{\text{Cl}^\bullet, 2\text{-Pr}} = 6.0 \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$ and $k_{\text{Cl}^\bullet, t\text{-But}} = 6.2 \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$) [54, 55].

4. Conclusion

In this paper UV fraction of solar light (UVA/B) is used to efficiently activate hydrogen peroxide and persulfate leading to the degradation of BPA, adopted as pollutant model, in simulated and real STPW. A strong effect of chloride ions is observed on the UVA/B– $\text{S}_2\text{O}_8^{2-}$ system where photogenerated sulfate radicals can lead to the formation of reactive species such as hydroxyl, chlorine and dichlorine radical anions and probably HOCl/OCl^- . In fact, at high chloride ions concentration an increase of BPA removal was observed.

In the presence H_2O_2 , carbonate ions strongly inhibit the efficiency of BPA degradation, while, this effect, is less pronounced using UVA/B– $\text{S}_2\text{O}_8^{2-}$. In STPW matrix the presence of Dissolved Organic Matter (DOM) represents the main radical scavenger; however, this negative effect can be counterbalanced by chloride ions using UVA/B– $\text{S}_2\text{O}_8^{2-}$.

These results are encouraging for future application of solar activation of persulfate for pollutant removal in STPW containing high chloride ions concentrations. Moreover, the use of chemical kinetic approach improves the oxidant capacity estimation of advanced oxidation process as function of water chemical composition.

Acknowledgements

This work was partially supported by Natural Science Foundation of China (No. 21367003) and Guangxi Natural Science Foundation (No. 2014GXNSFBA118217). We also gratefully acknowledge the Haina Project of Guangxi University for providing financial support for W.H. to stay at the Clermont Auvergne University in Clermont-Ferrand, France. Authors acknowledge financial support from the Region Council of Auvergne, from the “Fédération des Recherches en Environnement” through the CPER “Environment” founded by the Region Auvergne, the French government, FEDER from the European Community and PRC program CNRS/NSFC n°270437.

References

- [1] X. He, G. Zhang, A.A. de la Cruz, K.E. O'Shea, D.D. Dionysiou, Degradation mechanism of cyanobacterial toxin cylindrospermopsin by hydroxyl radicals in homogeneous UV/H₂O₂ process, *Environ. Sci. Technol.*, 48 (2014) 4495-4504.
- [2] J. Saien, M. Osali, A.R. Soleymani, UV/persulfate and UV/hydrogen peroxide processes for the treatment of salicylic acid: effect of operating parameters, kinetic, and energy consumption, *Desalin. Water Treat.*, 56 (2015) 3087-3095.
- [3] P. Xie, J. Ma, W. Liu, J. Zou, S. Yue, X. Li, M.R. Wiesner, J. Fang, Removal of 2-MIB and geosmin using UV/persulfate: Contributions of hydroxyl and sulfate radicals, *Water Res.*, 69 (2015) 223-233.
- [4] W. Yang, H. Zhou, N. Cicek, Treatment of organic micropollutants in water and wastewater by UV-based processes: A literature review, *Crit. Rev. Env. Sci. Tec.*, 44 (2014) 1443-1476.
- [5] S.-H. Yoon, S. Jeong, S. Lee, Oxidation of bisphenol A by UV/S₂O₈²⁻: Comparison with UV/H₂O₂, *Environ. Technol.*, 33 (2012) 123-128.
- [6] S.E. Beck, H. Ryu, L.A. Boczek, J.L. Cashdollar, K.M. Jeanis, J.S. Rosenblum, O.R. Lawal, K.G. Linden, Evaluating UV-C LED disinfection performance and investigating potential dual-wavelength synergy, *Water Res.*, 109 (2017) 207-216.
- [7] Y. Wu, A. Bianco, M. Brigante, W. Dong, P. de Sainte-Claire, K. Hanna, G. Mailhot, Sulfate radical photogeneration using Fe-EDDS: Influence of critical parameters and naturally occurring scavengers, *Environ. Sci. Technol.*, 49 (2015) 14343-14349.
- [8] B.M. Souza, M.W.C. Dezotti, R.A.R. Boaventura, V.J.P. Vilar, Intensification of a solar photo-Fenton reaction at near neutral pH with ferrioxalate complexes: A case study on diclofenac removal from aqueous solutions, *Chem. Eng. J.*, 256 (2014) 448-457.
- [9] L. Zhao, Y. Ji, D. Kong, J. Lu, Q. Zhou, X. Yin, Simultaneous removal of bisphenol A and phosphate in zero-valent iron activated persulfate oxidation process, *Chem. Eng. J.*, 303 (2016) 458-466.
- [10] C. Zhu, G. Fang, D.D. Dionysiou, C. Liu, J. Gao, W. Qin, D. Zhou, Efficient transformation of DDTs with Persulfate Activation by Zero-valent Iron Nanoparticles: A Mechanistic Study, *J. Hazard. Mater.*, 316 (2016) 232-241.
- [11] D. Xia, R. Yin, J. Sun, T. An, G. Li, W. Wang, H. Zhao, P.K. Wong, Natural magnetic pyrrhotite as a high-Efficient persulfate activator for micropollutants degradation: Radicals identification and toxicity evaluation, *J. Hazard. Mater.*, 340 (2017) 435-444.
- [12] P. Polczynski, R. Jurczakowski, W. Grochala, Stabilization and strong oxidizing properties of Ag(ii) in a fluorine-free solvent, *Chem. Commun.*, 49 (2013) 7480-7482.
- [13] J. Wang, S. Wang, Activation of persulfate (PS) and peroxymonosulfate (PMS) and application for the degradation of emerging contaminants, *Chem. Eng. J.*, 334 (2018) 1502-1517.
- [14] L.R. Bennedsen, J. Muff, E.G. Sogaard, Influence of chloride and carbonates on the reactivity of activated persulfate, *Chemosphere*, 86 (2012) 1092-1097.
- [15] Y. Liu, X. He, X. Duan, Y. Fu, D. Fatta-Kassinos, D.D. Dionysiou, Significant role of UV and carbonate radical on the degradation of oxytetracycline in UV-AOPs: Kinetics and mechanism, *Water Res.*, 95 (2016) 195-204.
- [16] Y. Yang, J.J. Pignatello, J. Ma, W.A. Mitch, Comparison of halide impacts on the efficiency of contaminant degradation by sulfate and hydroxyl radical-based advanced oxidation processes (AOPs), *Environ. Sci. Technol.*, 48 (2014) 2344-2351.

- [17] A. Afzal, P. Drzewicz, J.W. Martin, M. Gamal El-Din, Decomposition of cyclohexanoic acid by the UV/H₂O₂ process under various conditions, *Sci. Total Environ.*, 426 (2012) 387-392.
- [18] C. Liang, Z.-S. Wang, N. Mohanty, Influences of carbonate and chloride ions on persulfate oxidation of trichloroethylene at 20 °C, *Sci. Total Environ.*, 370 (2006) 271-277.
- [19] K. Mopper, X. Zhou, Hydroxyl radical photoproduction in the sea and its potential impact on marine processes, *Science*, 250 (1990) 661.
- [20] R.E. Huie, C.L. Clifton, Temperature dependence of the rate constants for reactions of the sulfate radical, SO₄⁻, with anions, *J. Phys. Chem.*, 94 (1990) 8561-8567.
- [21] R.E. Huie, C.L. Clifton, P. Neta, Electron transfer reaction rates and equilibria of the carbonate and sulfate radical anions, *Int. J. Radiat. Appl. Instrum. C Radiat. Phys. Chem.*, 38 (1991) 477-481.
- [22] J. Kiwi, A. Lopez, V. Nadtochenko, Mechanism and kinetics of the OH-radical intervention during Fenton oxidation in the presence of a significant amount of radical scavenger (Cl⁻), *Environ. Sci. Technol.*, 34 (2000) 2162-2168.
- [23] J.E.F. Moraes, F.H. Quina, C.A.O. Nascimento, D.N. Silva, O. Chiavone-Filho, Treatment of saline wastewater contaminated with hydrocarbons by the photo-Fenton process, *Environ. Sci. Technol.*, 38 (2004) 1183-1187.
- [24] Y. Yang, J.J. Pignatello, J. Ma, W.A. Mitch, Effect of matrix components on UV/H₂O₂ and UV/S₂O₈²⁻ advanced oxidation processes for trace organic degradation in reverse osmosis brines from municipal wastewater reuse facilities, *Water Res.*, 89 (2016) 192-200.
- [25] G.-D. Fang, D.D. Dionysiou, Y. Wang, S.R. Al-Abed, D.-M. Zhou, Sulfate radical-based degradation of polychlorinated biphenyls: Effects of chloride ion and reaction kinetics, *J. Hazard. Mater.*, 227-228 (2012) 394-401.
- [26] G.P. Anipsitakis, D.D. Dionysiou, M.A. Gonzalez, Cobalt-mediated activation of peroxymonosulfate and sulfate radical attack on phenolic compounds. Implications of chloride ions, *Environ. Sci. Technol.*, 40 (2006) 1000-1007.
- [27] J. Huang, S.A. Mabury, Steady-state concentrations of carbonate radicals in field waters, *Environ. Toxicol. Chem.*, 19 (2000) 2181-2188.
- [28] R. Zhang, P. Sun, T.H. Boyer, L. Zhao, C.-H. Huang, Degradation of pharmaceuticals and metabolite in synthetic human urine by UV, UV/H₂O₂, and UV/PDS, *Environ. Sci. Technol.*, 49 (2015) 3056-3066.
- [29] C. Wu, K.G. Linden, Phototransformation of selected organophosphorus pesticides: roles of hydroxyl and carbonate radicals, *Water Res.*, 44 (2010) 3585-3594.
- [30] O.S. Keen, K.G. Linden, Degradation of antibiotic activity during UV/H₂O₂ advanced oxidation and photolysis in wastewater effluent, *Environ. Sci. Technol.*, 47 (2013) 13020-13030.
- [31] Y. Lee, D. Gerrity, M. Lee, S. Gamage, A. Pisarenko, R.A. Trenholm, S. Canonica, S.A. Snyder, U. Von Gunten, Organic contaminant abatement in reclaimed water by UV/H₂O₂ and a combined process consisting of O₃/H₂O₂ followed by UV/H₂O₂: Prediction of abatement efficiency, energy consumption, and byproduct formation, *Environ. Sci. Technol.*, 50 (2016) 3809-3819.
- [32] F.L. Rosario-Ortiz, E.C. Wert, S.A. Snyder, Evaluation of UV/H₂O₂ treatment for the oxidation of pharmaceuticals in wastewater, *Water Res.*, 44 (2010) 1440-1448.
- [33] H.-W. Yu, T. Anumol, M. Park, I. Pepper, J. Scheideler, S.A. Snyder, On-line sensor monitoring for chemical contaminant attenuation during UV/H₂O₂ advanced oxidation process, *Water Res.*, 81 (2015) 250-260.
- [34] R. Zhang, Y. Yang, C.-H. Huang, N. Li, H. Liu, L. Zhao, P. Sun, UV/H₂O₂ and UV/PDS treatment of trimethoprim and sulfamethoxazole in synthetic human urine: Transformation products and toxicity, *Environ. Sci. Technol.*, 50 (2016) 2573-2583.

- [35] E. Kattel, M. Trapido, N. Dulova, Oxidative degradation of emerging micropollutant acesulfame in aqueous matrices by UVA-induced $\text{H}_2\text{O}_2/\text{Fe}^{2+}$ and $\text{S}_2\text{O}_8^{2-}/\text{Fe}^{2+}$ processes, *Chemosphere*, 171 (2017) 528-536.
- [36] M. Brigante, T. Charbouillot, D. Vione, G. Mailhot, Photochemistry of 1-Nitronaphthalene: A potential source of singlet oxygen and radical species in atmospheric waters, *J. Phys. Chem. A*, 114 (2010) 2830-2836.
- [37] E. Hayon, A. Treinin, J. Wilf, Electronic spectra, photochemistry, and autoxidation mechanism of the sulfite-bisulfite-pyrosulfite systems. SO_2^- , SO_3^- , SO_4^- , and SO_5^- radicals, *J. Am. Chem. Soc.*, 94 (1972) 47-57.
- [38] Y. Wu, R. Prulho, M. Brigante, W. Dong, K. Hanna, G. Mailhot, Activation of persulfate by Fe(III) species: Implications for 4-tert-butylphenol degradation, *J. Hazard. Mater.*, 322, Part B (2017) 380-386.
- [39] P. Wardman, Reduction potentials of one - electron couples involving free radicals in aqueous solution, *J. Phys. Chem. Ref. Data*, 18 (1989) 1637-1755.
- [40] P. Neta, R.E. Huie, A.B. Ross, Rate constants for reactions of inorganic radicals in aqueous solution, *J. Phys. Chem. Ref. Data*, 17 (1988) 1027-1284.
- [41] P. Maruthamuthu, P. Neta, Phosphate radicals. Spectra, acid-base equilibriums, and reactions with inorganic compounds, *J. Phys. Chem. A*, 82 (1978) 710-713.
- [42] Z.B. Alfassi, S. Mosseri, P. Neta, Reactivities of chlorine atoms and peroxy radicals formed in the radiolysis of dichloromethane, *J. Phys. Chem.*, 93 (1989) 1380-1385.
- [43] R. Yuan, S.N. Ramjaun, Z. Wang, J. Liu, Effects of chloride ion on degradation of Acid Orange 7 by sulfate radical-based advanced oxidation process: Implications for formation of chlorinated aromatic compounds, *J. Hazard. Mater.*, 196 (2011) 173-179.
- [44] M.J. Watts, K.G. Linden, Chlorine photolysis and subsequent OH radical production during UV treatment of chlorinated water, *Water Res.*, 41 (2007) 2871-2878.
- [45] M. Deborde, U. von Gunten, Reactions of chlorine with inorganic and organic compounds during water treatment—Kinetics and mechanisms: A critical review, *Water Res.*, 42 (2008) 13-51.
- [46] G.V. Buxton, C.L. Greenstock, W.P. Helman, A.B. Ross, Critical review of rate constants for reactions of hydrated electrons, hydrogen atoms and hydroxyl radicals (OH/O^-) in aqueous solution, *J. Phys. Chem. Ref. Data*, 17 (1988) 513-886.
- [47] J.S. Moore, G.O. Phillips, A. Sosnowski, Reaction of the carbonate radical anion with substituted phenols, *Int. J. Radiat. Biol. Relat. Stud. Phys. Chem. Med.*, 31 (1977) 603-605.
- [48] J. Sharma, I.M. Mishra, D.D. Dionysiou, V. Kumar, Oxidative removal of Bisphenol A by UV-C/peroxymonosulfate (PMS): Kinetics, influence of co-existing chemicals and degradation pathway, *Chem. Eng. J.*, 276 (2015) 193-204.
- [49] W. Huang, M. Luo, C. Wei, Y. Wang, K. Hanna, G. Mailhot, Enhanced heterogeneous photo-Fenton process modified by magnetite and EDDS: BPA degradation, *Environ. Sci. Poll. Res.*, 24 (2017) 10421-10429.
- [50] H. Katsumata, S. Kawabe, S. Kaneco, T. Suzuki, K. Ohta, Degradation of bisphenol A in water by the photo-Fenton reaction, *J. Photochem. Photobiol.*, A, 162 (2004) 297-305.
- [51] E.J. Rosenfeldt, K.G. Linden, Degradation of Endocrine Disrupting Chemicals Bisphenol A, Ethinyl Estradiol, and Estradiol during UV Photolysis and Advanced Oxidation Processes, *Environ. Sci. Technol.*, 38 (2004) 5476-5483.
- [52] A. Zafra, M. del Olmo, B. Suárez, E. Hontoria, A. Navalón, J.L.s. Vílchez, Gas chromatographic-mass spectrometric method for the determination of bisphenol A and its chlorinated derivatives in urban wastewater, *Water Res.*, 37 (2003) 735-742.
- [53] C.L. Clifton, R.E. Huie, Rate constants for hydrogen abstraction reactions of the sulfate radical, SO_4^- . Alcohols, *Int.*

J. Chem. Kinet., 21 (1989) 677-687.

[54] R. Mertens, C. von Sonntag, Photolysis ($\lambda = 254$ nm) of tetrachloroethene in aqueous solutions, J. Photochem. Photobiol., A, 85 (1995) 1-9.

[55] S. Takashi, M. Kazuhiro, H. Hiroyuki, K. Meiseki, On the reactivity of chlorine atoms towards alcohols, Chem. Lett., 16 (1987) 1429-1430.

Table 1 : pH, main anions concentrations and TOC determined for STPW used in this work.

pH	[Cl ⁻] (mM)	[HCO ₃ ⁻]/[CO ₃ ²⁻] (mM)	[SO ₄ ²⁻] (μM)	[NO ₃ ⁻] (μM)	[TOC] (mgC L ⁻¹)
7.9	2.58	4.16 (50 mgC L ⁻¹)	54	12.3	27.2

Table 2 : Second order rate constant determined for BPA and STPW before and after carbonates removal. STPW indicated Sewage Treatment Plan water without chemical modification while STPW* is STPW after removal of carbonates (see materials and methods). Second order rate constants are normalized for the number of all carbons (organic and inorganic). Errors are estimated to $\pm 3\sigma$, which was obtained from the scattering of the experimental data from the fitting line

	$k_{HO^{\bullet}}$	$k_{SO_4^{\bullet-}}$	$k_{Cl_2^{\bullet-}}$	$k_{CO_3^{\bullet-}}$
BPA	$(8.6 \pm 1.0) \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$	$(4.7 \pm 1.1) \times 10^9 \text{ M}^{-1} \text{ s}^{-1}$	$(8.3 \pm 0.8) \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$	$(9.2 \pm 1.0) \times 10^7 \text{ M}^{-1} \text{ s}^{-1}$
STPW	$(2.9 \pm 0.4) \times 10^3 \text{ L mgC}^{-1} \text{ s}^{-1}$	$(1.3 \pm 0.2) \times 10^4 \text{ L mgC}^{-1} \text{ s}^{-1}$		
STPW*	$(6.1 \pm 1.1) \times 10^3 \text{ L mgC}^{-1} \text{ s}^{-1}$	$(4.1 \pm 0.3) \times 10^4 \text{ L mgC}^{-1} \text{ s}^{-1}$		

Table 3. Percentage of photogenerated radicals reactivity in STPW and STPW*, on hydrogenocarbonates and on BPA estimated considering the pseudo-first order decay determined as products of the second order rate constant (obtained in this work or from literature) and initial concentrations.

		HO[•]		SO₄^{•-}	
STPW		98.4 %		99.8 %	
STPW*	HCO₃⁻	81.5%	16.6 %	98.9 %	~1 %
BPA		1.6 %		0.2 %	

Figures Captions

Figure 1: BPA degradation in the presence of H_2O_2 (2 mM) and $\text{S}_2\text{O}_8^{2-}$ (2 mM) in buffered and unbuffered solutions at pH 8.0 under UVA/B radiation.

Figure 2 Degradation profile of BPA ($[\text{BPA}]/[\text{BPA}]_0$) at pH 4.6 (full symbols) and 8.0 (empty symbols) using different chloride ions concentrations.

Figure 3 BPA degradation efficiency (DE) using different chloride ions concentrations at pH 8.0 compared to the value obtained using 2 mM of persulfate alone. Insert: effect of chloride ions using a linear scale for chloride ion concentration.

Figure 4: Modeled radical concentration in the system UVA/B- $\text{S}_2\text{O}_8^{2-}$ (2 mM) in the presence of different chloride ions concentrations.

Figure 5: Inhibition effect of carbonates as a function of initial concentration and pH of solution toward the reactivity of hydroxyl (**A**) and sulfate (**B**) radicals on BPA. An initial concentration of 43 μM of BPA is considered.

Figure 6: BPA 430 nM (0.1 mg L^{-1}) degradation in STPW using UVA/B- H_2O_2 (2 mM) and UVA/B- $\text{S}_2\text{O}_8^{2-}$ (2 mM) system. STPW indicates sewage treatment plant water without any chemical modification; STPW* indicates sewage treatment plant water after removal of carbonates as indicated.

Figure 7: 4.3 μM BPA degradation in STPW using $\text{S}_2\text{O}_8^{2-}$ (2 mM) under UVA/B irradiation in the

presence of 50 mM of 2-propanol (2-Pr) and 50 mM of *tert*-Butanol (*t*-But).

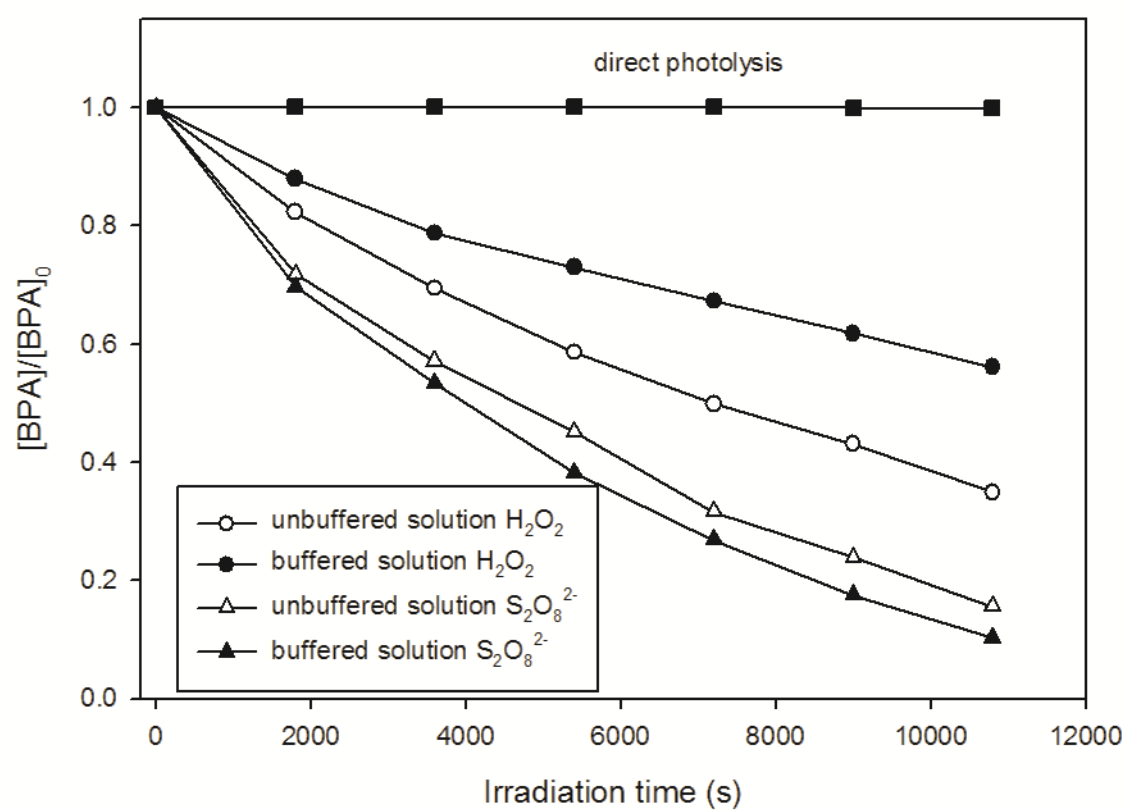
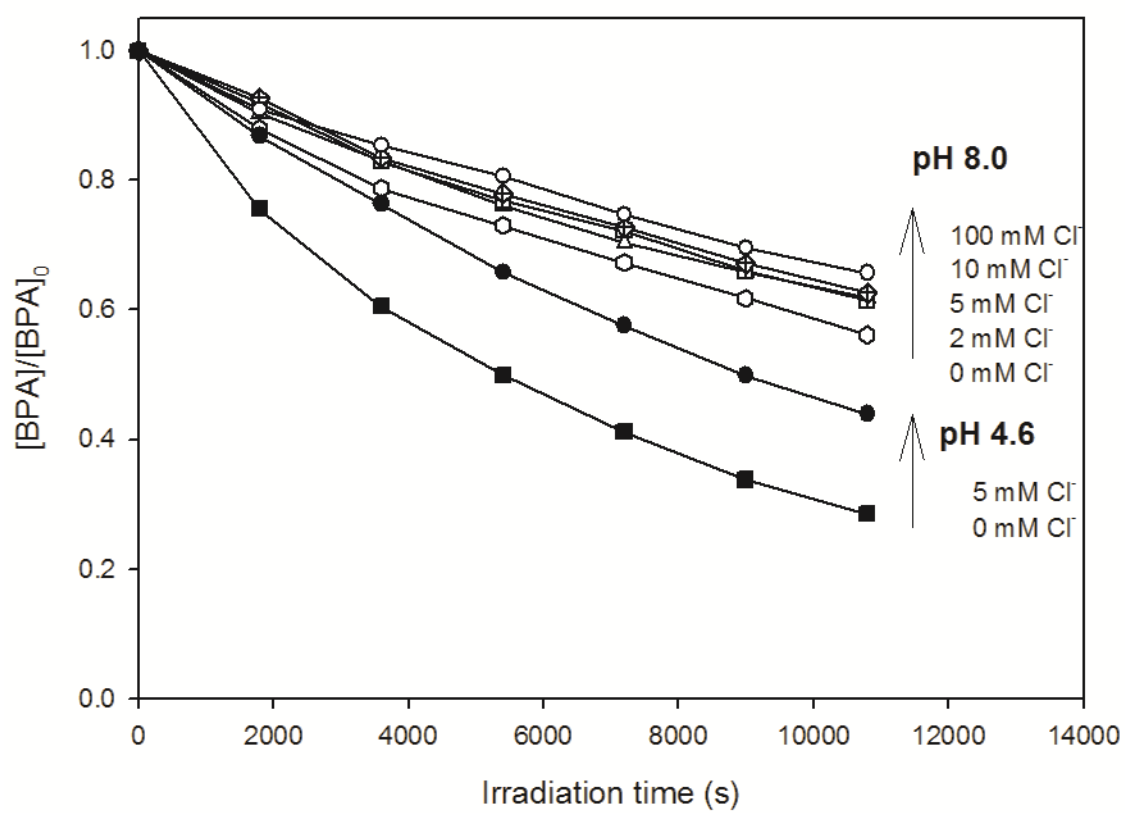
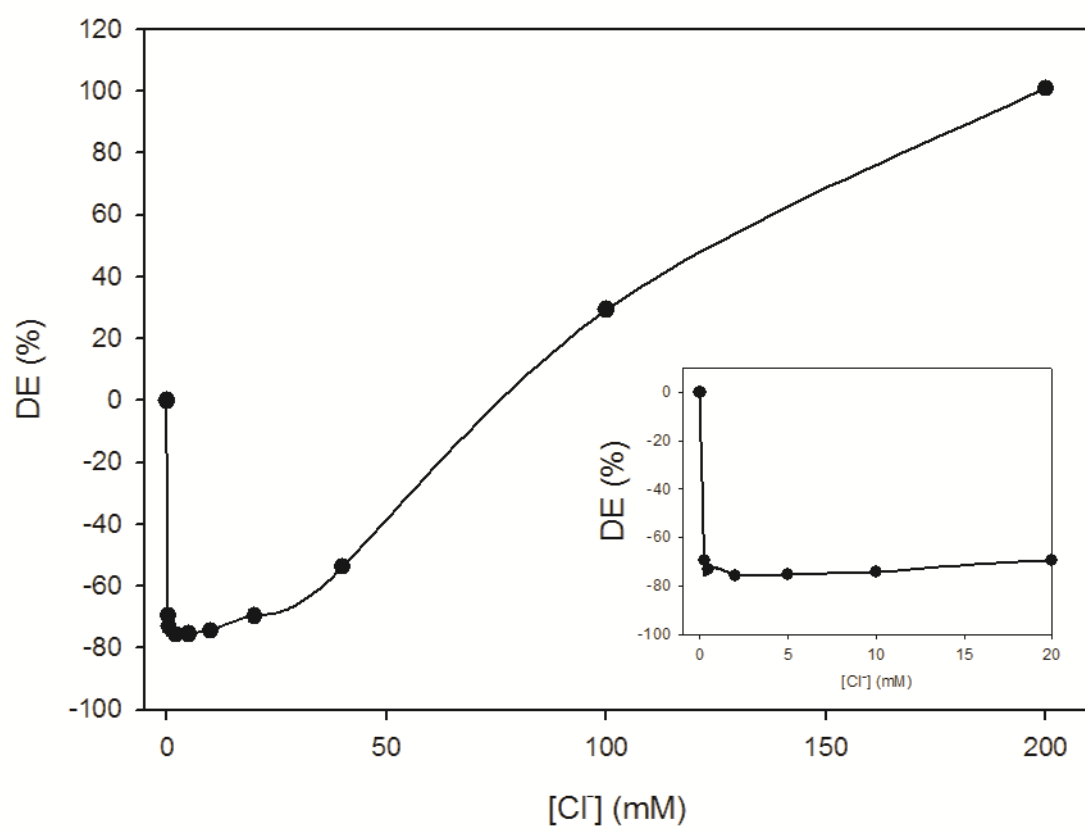
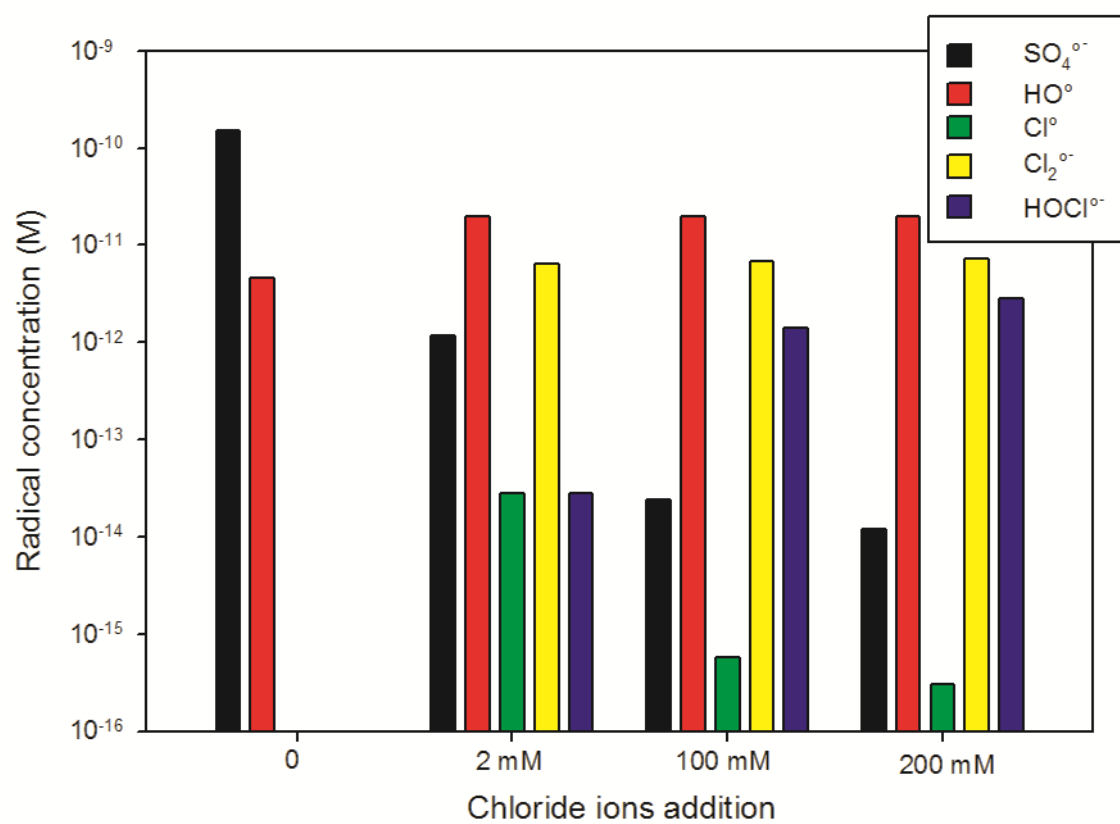
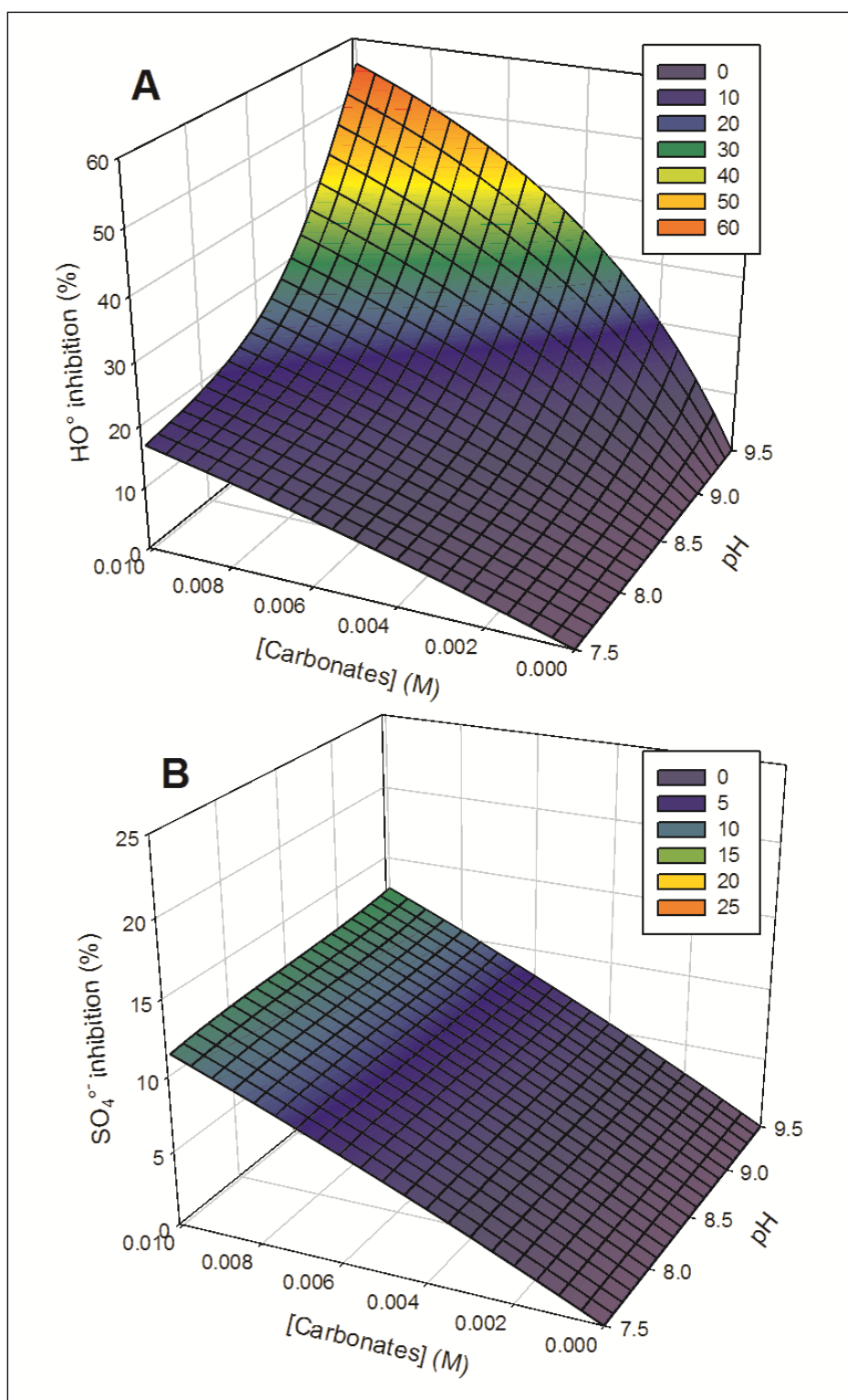


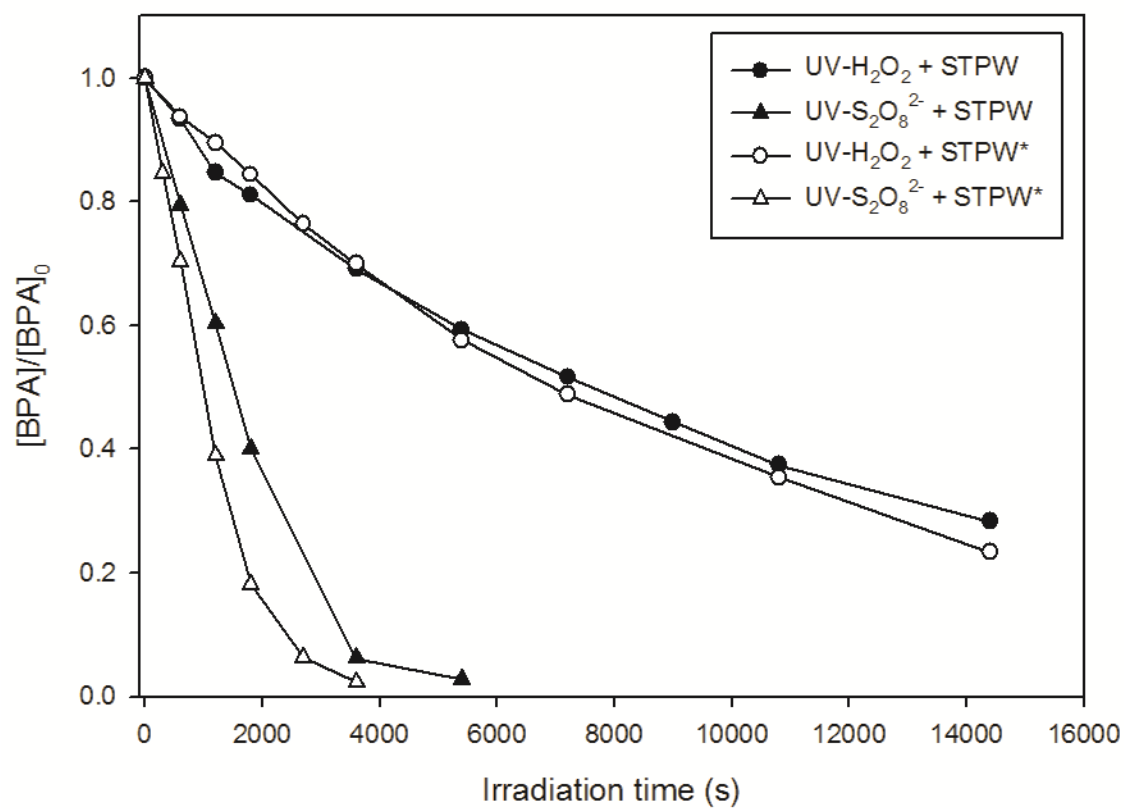
Figure 1

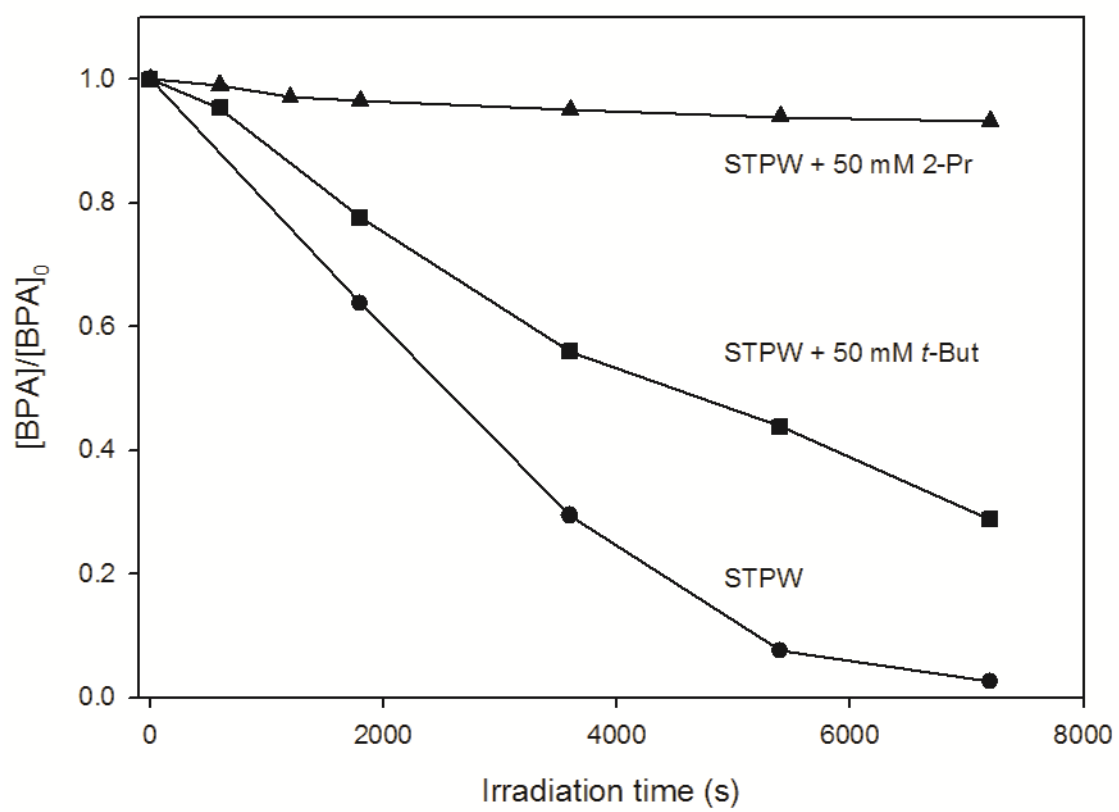
**Figure 2**

**Figure 3**

**Figure 4**

**Figure 5**

**Figure 6**

**Figure 7**