Study on anodic oxidation parameters for removal of pesticide imidacloprid on a modified tantalum surface by lead dioxide film

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Abstract

The commercial imidacloprid (IMD) insecticide [1-(6-chloro-3-pyridinyl) methyl-4,5-dihydro-N-nitro-1H-imidazole-2-amine] is widely used for the enhancement of crop production, but the intensive use of this insecticide has caused serious environmental problems. This work presents an electrochemical process for the removal of this insecticide using galvanostatic electrolysis at modified tantalum surface by lead dioxide film anode (Ta(PbO$_2$)) anode. The electrolytic process was monitored by chemical oxygen demand (COD). The influence of operating parameters, such as current density, initial concentration of IMD, temperature and initial pH value was investigated. The COD decay follows a pseudo first-order kinetic and the process was under mass transport control. COD removal reach 97% when using an apparent current density of 100 mA cm^{-2}, initial COD of 953 mg L^{-1} and at 25 °C after 4.5 h electrolysis time.

Keywords: Imidacloprid, electrochemical degradation, anodic oxidation, Ta(PbO$_2$) anode, hydroxyl radicals.

1. Introduction

Crop yield is often minimized by the presence of undesirable plants, insects and various diseases. For each of these "enemies", the industry has created a defense commonly known pesticide. However, the intensive use of these pesticides in agriculture leads to adverse impacts in the aquatic and terrestrial ecosystems. In addition, several studies have found that, many pesticides are more toxic and environmentally persistent, creating serious acute health problems and environmental impacts (Ecobichon, 2012). Many studies showed that pesticides were detected i*n agricultural* products (Wen *et al.*, 2013; Paranthaman *et al.*, 2012) and in groundwater (Johnson *et al.*, 2001; Kadian *et al.*, 2008).

Imidacloprid [1-(6-chloro-3-pyridylmethyl)-N-nitroimidazolidin-2-ylidenamine] a relatively stable chloronicotinic insecticide, classified as Category I due to its high leaching potential (USEPA, 2009). It could easily contaminate water sources, leading to negative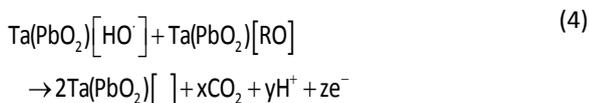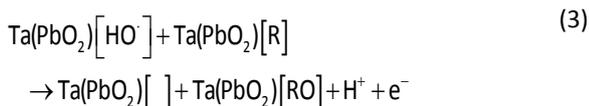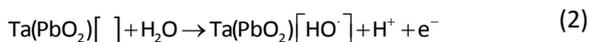 environmental and health effects. Indeed, acute exposure to Imidacloprid may cause apathy, spasms and thyroid lesions. This insecticide is used to control a number of agricultural insect such as aphids, whiteflies, leafhoppers, termites (Mullins, 1993; Tomizawa and Casida, 2005). IMD residues may occur in water and agricultural products therefore they pose a potential hazard for consumers. For these reasons, it is necessary to develop technologies for the removal of this compound.

Advanced Oxidation Process (AOP) is a chemical treatment method that has been growing in the wastewater management industry. It is a highly suggested method for the removal of problematic organic matter. The basic principle of AOP involves the production of hydroxyl radicals (HO$^•$), which can be generated from hydrogen peroxide (H$_2$O$_2$), ozone, photo-catalysis, or oxidants in combination with using ultraviolet (UV) radiation. In some cases, two or more radical generators are used in combination. However, it is the HO$^•$ that is mainly responsible for the degradation of organic compounds. This radical then goes through a succession of chemical reactions to form several products and by-products (Bokare and Choi, 2014). Theoretically, AOPs should fully mineralize organic compounds to carbon dioxide and water as illustrated in equation 1.

$$RH + HO^• \rightarrow {}^•R + H_2O \qquad (1)$$

Anodic oxidation or electro-oxidation is one of the most popular advanced oxidation procedures for elimination of dissolved organic pollutants and consists in the oxidation of organic matter in an electrolytic cell. In anodic oxidation, organic pollutants are directly destroyed by reaction with hydroxyl radical (HO$^•$) formed at the anode surface from water oxidation (Torres *et al.*, 2003).

When using Ta(PbO$_2$) anode, properties of this material enable the oxidation of organic compounds mediated by hydroxyl radicals. The mechanism consists in the H$_2$O oxidation at the anode surface to produce a slightly adsorbed hydroxyl radical on the electrode that reacts with the organic molecules (R) that reach the vicinity of the anode producing CO$_2$ and H$_2$O as shown in the following equations:

$$Ta(PbO_2)[\ ] + H_2O \rightarrow Ta(PbO_2)[HO^-] + H^+ + e^- \qquad (2)$$

$$Ta(PbO_2)[HO^-] + Ta(PbO_2)[R] \qquad\qquad (3)$$
$$\rightarrow Ta(PbO_2)[\ ] + Ta(PbO_2)[RO] + H^+ + e^-$$

$$Ta(PbO_2)[HO^-] + Ta(PbO_2)[RO] \qquad\qquad (4)$$
$$\rightarrow 2Ta(PbO_2)[\ ] + xCO_2 + yH^+ + ze^-$$

Thus, to oxidize such effluent, anode material should have besides high overvoltage for oxygen evolution, high electrical conductivity, chemical inertness and low cost. Among the possible electrode materials which meet these criteria, the so-called dimensionally stable anodes (DSA) (e.g., PbO_2 and SnO_2) have received considerable attention in recent years. These anodes have been successfully employed to destroy a variety of pollutants, phenols (Xiaoyue *et al.*, 2013), dyes (José *et al.*, 2013), pesticides (Olfa *et al.*, 2013) and so on.

The goal of the present work was attempted to use electrocatalytic oxidation using $Ta(PbO_2)$ anode for the treatment of IMD in aqueous solutions. The electrocatalytic oxidation of IMD was followed by chemical oxygen demand, UV-Vis spectrophotometric and square wave voltammetry method.

2. Experimental

2.1. Reagents

Solutions contain quantities of IMD taken from an emulsifiable concentrate (Confidor 200 SL from Bayer CropScience) containing 200 g L^{-1} IMD. All solutions were freshly prepared with ultrapure water (modelMillipore Direct-Q™ 3). Sodium hydroxide (NaOH) (0.005 M) and sulfuric acid (H_2SO_4) (0.01 M) of analytic grade were employed as conductive electrolytes and for pH adjustment.

2.2. Preparation of the modified tantalum surface

2.2.1. Tantalum surface treatment

Pre-treatments of the tantalum substrate (rectangular tantalum plates 5 cm × 1 cm × 1 mm) were carried out before anodization to ensure good adhesive lead dioxide film. Tantalum was first roughened to increase the adhesion of PbO_2 deposit via subjecting its surface to mechanical abrasion using silica grains with an average diameter of 0.3 mm projected under 5 bar pressure. It was then cleaned to remove sand particles or any other particles lodged in the metal surface. This was carried out by degreasing with acetone because of its ease of application and its great penetrating power, and then ultrasonically rinsed in ultrapure water during 10 min. Uniform and well adhesive deposit necessitates a smooth surface with no oxides (formed spontaneously on contact with oxygen in the air) or scales. To ensure this, the tantalum substrate was soaked for 30 s in hydrofluoric acid (40% weight) at room temperature and then abundantly rinsed with ultrapure water. This chemical treatment corresponds to dissolution of the surface layer formed of tantalum and tantalum oxide according to following reactions:

$$Ta + 5HF \rightarrow TaF_5 + \frac{5}{2}H_2 \qquad\qquad (5)$$

$$Ta_2O_5 + 10HF \rightarrow 2TaF_5 + 5H_2O \qquad\qquad (6)$$
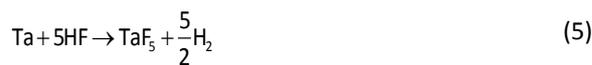
2.2.2. Electrochemical deposition of PbO_2 film

The lead dioxide was deposited galvanostatically on the pretreated tantalum substrate by electrochemical anodization of an aqueous $Pb(NO_3)_2$ solution (1 M) placed in a two-compartment Pyrex glass cell (V = 150 cm^3) thermoregulated at 65 °C. The electrodeposition of PbO_2 film was carried out at an apparent current density of 10 mA cm^{-2} for 0.5 hours, then at 20 mA cm^{-2} for the same period and finally at 50 mA cm^{-2} for one hour. The average masse of PbO_2 was 0.21 g cm^{-2}. The deposit obtained was a grey porous material with strong adherence.

2.3. Electrode characterization

The morphology of the electrode was examined using an optical microscope (SZ-45-PS; CALTEX Scientific Inc., Irvine, CA).

The X-ray diffraction (XRD) patterns were recorded on a D8 Advance X-diffractometer (Cu K_α, λ = 0.154 nm) (Bruker, Germany) operating at 40 kV and 30 mA. The diffraction patterns were collected at 25 °C and over an angular range of 5 to 100° with a step size of 0.05° per step and a dwell time of 12 s per increment.

2.4. Electrolysis

Galvanostatic electrolysis of IMD aqueous solutions (150 cm^{-3}) was carried out in a two-compartment thermostatic cell (Figure 1). The cathode is a cylindrical platinum grid (φ = 4 cm, l = 5 cm). The anode is made up of $Ta(PbO_2)$ anode plate arranged symmetrically around the cathode. The geometric working surface of each plate was 5 cm^2. IMD solutions were electrolyzed in galvanostatic mode using a DC power supply (model ABTP 530 Françoise d'Instrumentation, France). The range of applied current density was 50 to 100 mA cm^{-2}. The pH of solution was adjusted, before and over the course of the electrolysis, by adding either concentrated sulfuric acid or sodium hydroxide solutions.
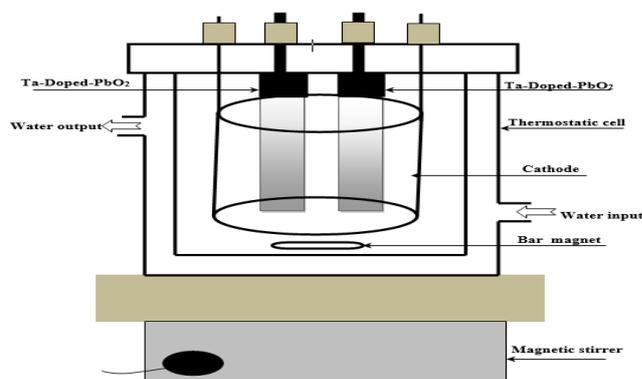

Figure 1. Electrolytic cell

2.5. Analysis

2.5.1. Chemical oxygen demand (COD)

The COD measurement during the processing permitted the evaluation of the kinetic of organic matter decay and the instantaneous current efficiency (ICE). COD was determined by the dichromate method (Kolthof *et al.*, 1969).

The absorbance measurements were carried out with a UV–visible spectrophotometer (Shimadzu 1650 PC).

2.5.2. Square-wave voltammetry analysis

The Square-wave voltammetry measurement was performed using a potentiostat-galvanostat (VoltaLab PST050), and a conventional three-electrode cell at 25 °C. BDD was used as working electrode for voltammetric experiments. A platinum wire was used as a counter electrode and a saturated calomel electrode (SCE) was used as reference electrode.

2.5.3. UV-Vis spectrophotometer analysis

The treated samples were analyzed using UV-Vis spectrophotometer (Shimadzu 1650 PC). The samples were filtered through Millipore filters (0.45 μm) and absorbance of the compound was measured at corresponding λmax at 270 nm. From standard solutions, calibration curves were prepared and concentrations of experimental solutions were thus determined.

3. Results and discussion

3.1. Crystalline structure of Ta(PbO₂) electrode

Ta(PbO_2) electrode is one of the most widely investigated anode materials for electrochemical oxidation due to its chemical stability, high electronic conductivity, and low cost of material (Chen, 2004). The XRD patterns of Ta(PbO_2) anode electrode are shown in Figure 2. The X-ray diffraction data were recorded by using Cu K_α radiation (1.54 Å). The intensity data were collected over a 2θ range of 5–100. As can be seen, several remarkable peaks at 23.63°, 29.09°, 32.30°, 38.09°, 49.53°, 63.63°, 76.36° and 80° are attributed to the β-PbO_2 substrate (Xiaoyue *et al.*, 2013; José *et al.*, 2013; Xu *et al.*, 2015).
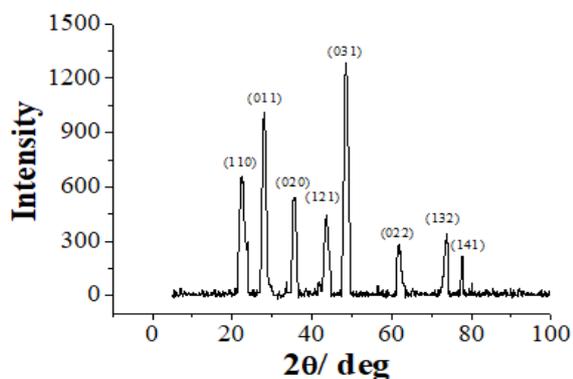

Figure 2. X-ray diffraction (XRD) pattern of β-PbO_2 electrode prepared by electro-deposition

3.2. Surface morphology

The morphology of the prepared electrodes was investigated by optical microscope as shown in Figure 3. Before the polishing and chemical stripping in hydrofluoric acid 40%, the tantalum substrate micrographs presented a rough and reproducible surface. However, after the electrochemical deposition film, the lead oxide was stable and adherent almost on all tantalum plate surfaces. Furthermore, the Ta(PbO_2) anode deposit is very porous and presents a high roughness.
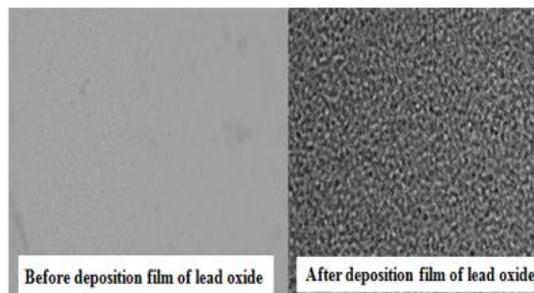

Figure 3. The morphology of the prepared of Ta(PbO_2) anode by optical microscope in electrolysis of IMD aqueous solution before and after chemical deposition of lead oxide

3.3. Stability of Ta(PbO₂) electrode

The dissolution of PbO_2 was studied by following changes in the concentration of lead in solution during the electrolysis of a solution of IMD during all experimental electrolysis on the Ta(PbO_2) anode. Figure 4 shows that, the lead oxide dissolves rapidly in the first hours of electrolyses, and then the concentration of lead in solution increases slightly with the electrical charge. Should be noted that. the standard in drinking water and surface water intended is 15 mg L^{-1} according to the new European directive (USEPA, 2009).

To test the repeatability of electro-oxidation efficiency of Ta(PbO_2) anode, 4 anodic oxidation tests were continued using a 500 mg L^{-1} aqueous solution of IMD, corresponding to a COD value of about 953 mg L^{-1}. After each test, the Ta(PbO_2) anode was only washed by distilled water and dried using a blower. After 4.5 h of each test of electrolysis the experimental results demonstrate a constant efficiency of COD removal in the range of 95–97%.
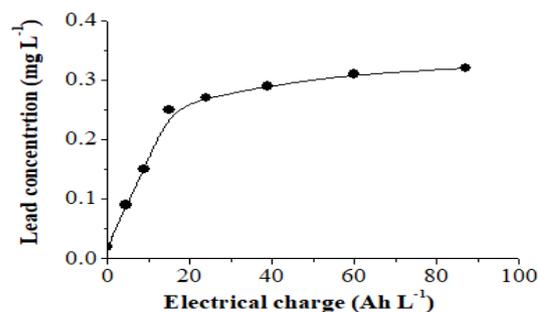

Figure 4. Evolution of lead concentration as a function of electrical charge measured during all tests done in this work

3.4. Influence of current density

The role of current density has been followed, testing values smaller that the previously considered as the optimum. It corresponds to the ratio between the applied current and the surface of the working electrode. Therefore, the current density can be altered by changing the current and/or the surface of the working electrode.

To achieve complete mineralize of IMD to CO_2 and H_2O, 66 mole of $HO^{\bullet}$ is necessary to mineralize one mole of IMD (Eq. (7))

$$C_9H_{10}ClN_5O_2 + 66HO^{\cdot} \rightarrow 9CO_2 + 35H_2O + 5HNO_3 + HCl \qquad (7)$$
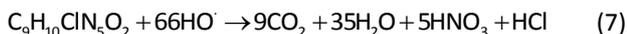
So, it is necessary to choose suitable current density for the generation of amount of hydroxyl radicals sufficient to accomplish mineralization of IMD to CO_2 and H_2O.

The influence of the applied current density on the COD removal during the electrochemical oxidation of IMD at the $Ta(PbO_2)$ anode is shown in Figure 5. It could be seen from this figure that the removal of COD significantly enhanced with the increase of current density. For example, after 270 minutes of electrolysis the COD percent removal increased from 24% to 97% when the current density increased from 50 to 100 mA cm^{-2}. The process could be attributed to an increase of the amount of hydroxyl radicals with the increase of the applied current (Belhadj Ammar *et al.*, 2016; Rkik *et al.*, 2017; Ben Brahim *et al.*, 2016). However, increasing of the current density beyond 100 mA cm^{-2} involves a reduction the yield of the degradation reaction. The oxidation rate decreased due to the rise of side reactions such as oxygen evolution (Eq. (8)) and/or electrolyte decomposition (Eq. (9)) which occur at the anode surface (Panizza and Cerisola, 2005; Sires *et al.*, 2008).

$$2H_2O \rightarrow O_2 + 4H^+ + 4e^- \qquad (8)$$

$$2SO_4^{2-} \rightarrow s_2O_8^{2-} + 2e^- \qquad (9)$$

Therefore, the current density of 100 mA cm^{-2} was selected as the optimal parameter in the following experiments.

The instantaneous current efficiency (ICE) can be defined as the art of the current directly used for the oxidation of the organic compounds (inset of Figure 5). The increased current density resulted in a decrease in current efficiency. The ICE can also be defined by Eq. (10) (Comninellis and Pulgarin, 1991).

$$ICE = \frac{COD(t) - COD(t + \Delta t)}{8It} FV \qquad (10)$$

where, F is the Faraday constant (96,487 C/mol), V the volume of the solution (L), COD(t) and COD ($t + \Delta t$) are the chemical oxygen demands at times t and t + Δt (in g O_2 L^{-1}), respectively, and I is the current (A).

We notice that the instantaneous current efficiency is below 100% (inset of Figure 5) revealing that the degradation process of IMD was controlled more by electrode kinetics than by mass transfer in the tested current density range.
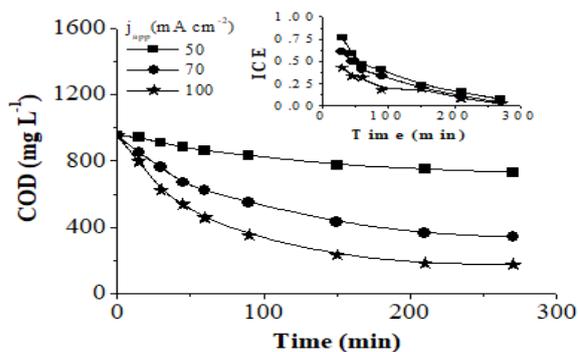

Figure 5. Influence of the apparent applied current density on the trends of COD and ICE during electrolysis of IMD (COD$_0$ 953 mg L^{-1}) on the Ta(PbO$_2$) anode. Electrolyte: Na$_2$SO$_4$ 2 g L^{-1}; pH = 3.0 and T = 25 °C

The decay of COD exhibits an exponential behavior with all applied current indicating a first-order reaction kinetics. If we suppose that the active surface of the electrode remains constant during electrolysis, we can consider that the hydroxyl radical's concentration is in a steady state and therefore, the oxidation rate expression can be written by Eq. (11):

$$r = -\frac{dCOD}{dt} = k[HO^{\cdot}]^{\alpha} COD(t) = k_{app} COD(t) \qquad (11)$$

where, α is the reaction order related to the hydroxyl radicals, k is the real rate constant, k_{app} is the global apparent rate constant for COD removal.

The integration of the previous equation leads to the Eq. (12):

$$COD(t) = COD(0) \exp k_{app} t \qquad (12)$$

where k_{app} could be calculated from the slope value of the plot of $\ln[COD(0)/COD(t)]$ versus t (Figure 6). The straight lines obtained in these plots were in agreement with a pseudo first-order COD removal.
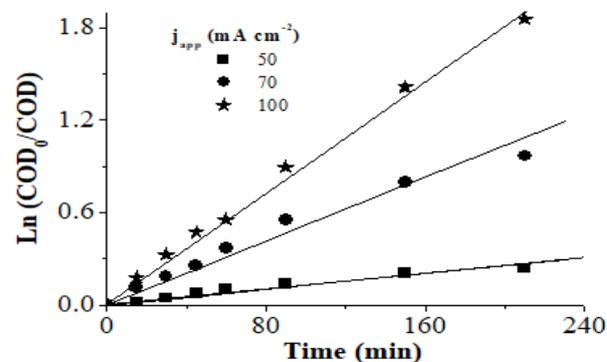

Figure 6. The linear regression for the COD removal with time during electrolysis of the IMD (COD$_0$ 953 mg L^{-1}) on the Ta(PbO$_2$) anode for different current densities. Electrolyte: Na$_2$SO$_4$ 2 g L^{-1}; pH = 3.0 and T = 25 °C

In addition, the surveillance of IMD concentration during the treatment by anodic oxidation process was followed by SWV at BDD electrode. The comparison between the COD removal and the IMD concentration during the electrolysis time is introduced in Figure 7. The results show that the molecules of IMD disappear faster than the overall organic material expressed in terms of COD. Indeed, the IMD molecules disappear completely after 200 minutes, but at the same time only 75% of the COD was removed, this explains by the formation of degradation products during the electro-oxidation of the IMD which are required more time to finish disappearing. A general reaction sequence for the electrochemical degradation of IMD is proposed in the Figure 8 (Yujing *et al.*, 2013).
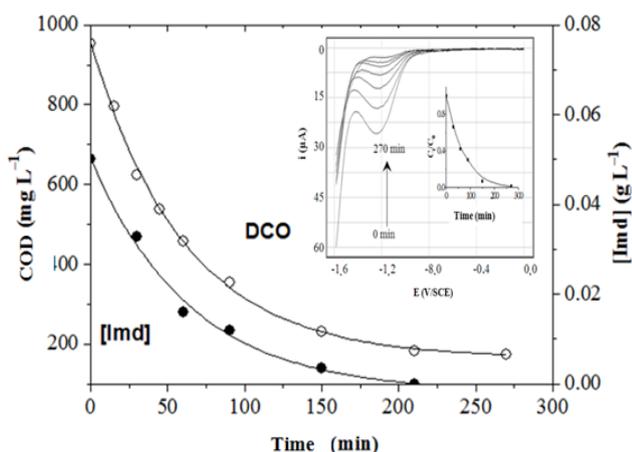

Figure 7. Evolution of COD removal and IMD concentration (followed by SWV) during electrolysis of IMD (COD_0 953 mg L^{-1}) on the Ta(PbO_2) anode. $f = 100$ s^{-1}, $\Delta E_i = 2$ mV, $\Delta E_a = 50$ mV, pH = 3 and T = 25 °C
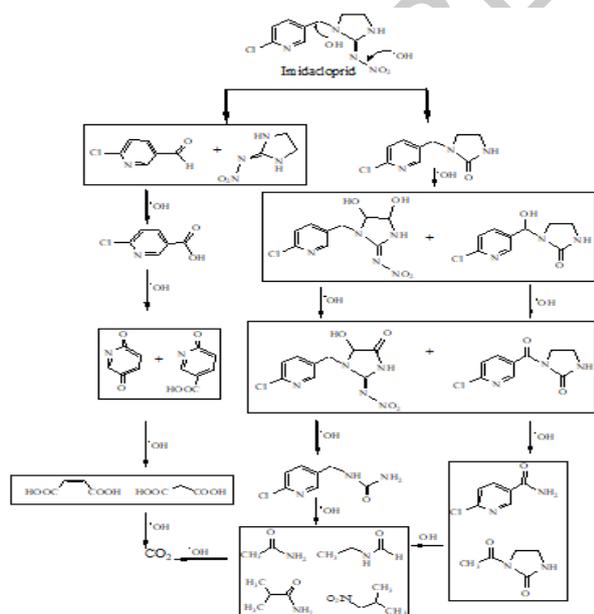

Figure 8. The proposed reaction sequence for the degradation of the IMD by ES-EF system (Yujing *et al.*, 2013)

3.5. Influence of pH

Solution pH is an important factor for wastewater treatment. Various works have reported on the effects of solution pH in anodic oxidation of organic compounds on PbO_2 anode, but the results are contradictory. For example, Kirk *et al.* (1985) found that the current efficiency for the oxidation of aniline at a PbO_2 anode increased from 3% to 13% when pH improved from 2 to 11 (Kirk *et al.*, 1985). In contrast, in a previous study found that the oxidation kinetics of 4-chloroguaiacol removal using PbO_2 anode was faster at pH 2 than at pH 6 (Samet *et al.*, 2006). On the other hand, Panizza *et al.* (2008) demonstrated that the effect of pH was not significant for the oxidation of methyl red at PbO_2 and BDD anodes (Panizza and Cerisola, 2008). The same result was observed by Chen and Chen (2006) when they studied the oxidation of orange II at a BDD anode (Chen and Chen, 2006). These discrepancies can be explained by the differences in the chemical properties of the organic compounds tested and the characteristic of the electrode materials (Cui *et al.*, 2017). To study the influence of pH in the oxidation of IMD, electrolysis was carried out in the pH range of 3–10 (Figure 9).
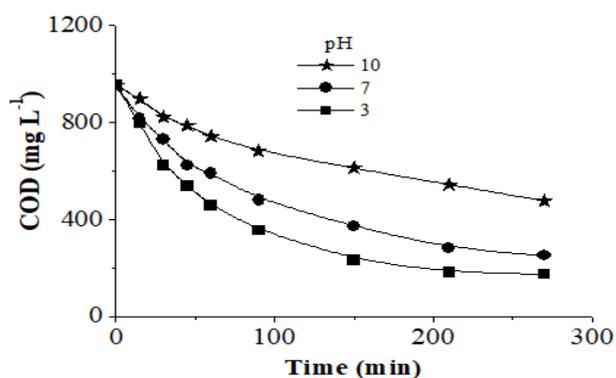

Figure 9. The influence of pH on the trends of COD during electrolysis of the IMD (COD_0 953 mg L^{-1}) on the Ta(PbO_2) anode. Electrolyte: Na_2SO_4 2 g L^{-1}; $j_{app} = 100$ mA cm^{-2} and T = 25 °C

As can be seen from this figure, the degradation of IMD in acid medium is more efficient than in alkaline and neutral medium. This effect may be due to the extensive oxidation and/or chemical modification of the electrode surface, which suggests a change in the surface properties. This result is in agreement with previous literature reports for degradation of some pesticides (El Mouden *et al.*, 2012). In pH = 10, we remark that the solutions become a little bit turbid after oxidation. This indicates the formation of polymeric intermediate products by the hydroxyl group that makes the degradation much more difficult (Massa *et al.*, 2017; Rabaaoui *et al.*, 2013). Therefore, a pH 3 was chosen as the optimal value.

3.6. Thermodynamics studies of IMD

In order to study the effect of reaction temperature on the removal of IMD, data were collected at four different temperatures (25, 35, 45 and 65 °C) at pH 3, $j_{app} = 100$ mA cm^{-1}, and $COD_0 = 953$ mg L^{-1}. The Figure 10 show that the COD removal was increased when the temperature increase from 25 to 65 °C. The result is

probably due to the increase in the indirect oxidation reaction of organics with $S_2O_8^{-2}$ from the analyte (Na_2SO_4) oxidation. This result is in agreement with those obtained by other authors (Serrano *et al.*, 2002).
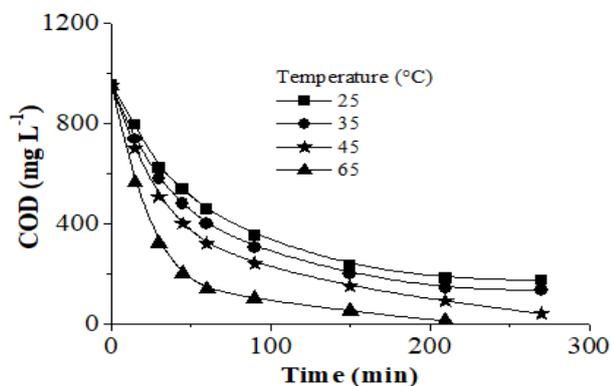
$$2SO_4^{2-} \rightarrow S_2O_8^{2-} + 2e^- \tag{13}$$

Figure 10. Influence of the temperature on the trends of COD during electrolysis of the IMD (COD_0 953 mg L^{-1}) on the Ta(PbO$_2$) anode. Electrolyte: Na_2SO_4 2 g L^{-1}; j_{app} = 100 mA cm^{-2} and pH = 3.0

Rate constants for chemical reactions are known to depend strongly on the reaction temperature. One well known empirical relationship expressing this dependence is the Arrhenius equation (14).

$$k_{app} = A \exp\left(-\frac{E_{app}}{RT}\right) \tag{14}$$

The activation free energy (ΔG*), the activation enthalpy (ΔH*) and the activation entropy (ΔS*) for the anodic oxidation process can be determined from the following equations:

$$\Delta H^* = E_a - RT \tag{15}$$

$$\Delta S^* = R\left(\ln A - \ln\frac{k_B T}{h} - 1\right) \tag{16}$$

$$\Delta G^* = \Delta H^* - T\Delta S^* \tag{17}$$

where k_{app} is the global apparent rate constant, A is the preexponential factor (s^{-1}), E_{app} is the activation energy (J mol^{-1}), k_B is the Boltzmann's constant, h is the Planck's constant, R is the ideal gas constant and T is the reaction absolute temperature.

Table 1. Thermodynamic parameters for anodic oxidation of IMD

T (°C)	E_a (kJ mol^{-1})	ΔH^* (kJ mol^{-1})	ΔS^* (J mol^{-1} k^{-1})	ΔG^* (kJ mol^{-1})
25	19.70	17.222	-8,05	19.620
35		17.139	-7.37	19.408
45		17.057	-6.169	19.018
65		16. 889	-4.087	18.270

The results of the thermodynamic calculations are shown in Table 1. The apparent activation energy is 19.70 kJ mol^{-1}. This value is less than 40 kJ mol^{-1} indicating that the system was under mass transport control (Belhadj-Tahar and Savall, 1998). Moreover, the positive value for the Gibbs free energy at all temperatures indicate that the oxidation reaction of IMD is non-spontaneous in nature and that the degree of the non-spontaneity of the reaction decreased by increasing temperature. The positive ΔH* refer to endothermic reaction and the negative value of ΔS* suggests that the intermediate formed during the oxidation of IMD is more ordered than the reactants.

4. Conclusion

This work is a first attempt to investigate the degradation of IMD in electrochemical treatment with Ta(PbO$_2$) anode. Electrochemical oxidation is a method that has been utilized to date for the treatment of several types of wastes. This paper provides new results on the electrochemical degradation of IMD and led to the following points:

• Ta(PbO$_2$) anode displayed noticeable oxidation abilities for treating wastewaters containing this pesticide, due to the production of a large amount of hydroxyl radicals on the electrode surface during electrolysis.

• the removal rate of COD increases with applied current density until 100 mA cm^{-2} due to the increase of the mass transport caused by oxygen evolution reaction,

• the evolution of COD with electrolysis time followed a pseudo first-order kinetics.

• the increased current density resulted in a decrease in current efficiency. Moreover, an increase in IMD concentration enhanced the degradation rate due to the increase in the organic matter diffusion flux as well as a decrease of the oxygen's formation rate.

• the effect of temperature showed that for 25 °C and 65 °C the achieved reduction was 82 % and 99 %. The apparent global activation energy was found as 19.70 kJ mol^{-1}.

• the increases of pH from acidic to alkaline medium decreased slightly the electrochemical removal of IMD.

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