

Electrocatalytic performance of PbO_2 films in the degradation of dimethoate insecticide

O Dridi Gargouri, Y Samet* and R Abdelhedi

UR Electrochimie et Environnement, Department of Materials Engineering, National Engineering School of Sfax, BPW 3038 Sfax, University of Sfax, Tunisia

Abstract

This study was performed to find the best experimental conditions for the electrochemical removal of the insecticide dimethoate ($\text{C}_3\text{H}_{12}\text{NO}_3\text{PS}_2$) from aqueous solutions using a lead dioxide niobium anode. The process was studied under galvanostatic polarisation mode. The influence of applied current density (10–50 $\text{mA}\cdot\text{cm}^{-2}$), initial chemical oxygen demand COD_0 (100–550 $\text{mg}\cdot\text{L}^{-1}$), temperature (30–70°C) and pH (3–11) on COD and instantaneous current efficiency (ICE) was studied. The results showed that almost 90% of COD removal was achieved under optimal experimental conditions, indicating that electrochemical oxidation on a PbO_2 anode is a suitable method for treatment of water polluted with dimethoate. It was found that the decay of COD generally followed a pseudo first-order kinetic and the oxidation rate was favoured by increasing the applied current density, temperature, pH and initial COD. The greatest COD removal (90%) was obtained when using an applied current density of 50 $\text{mA}\cdot\text{cm}^{-2}$, $\text{COD}_0 = 320 \text{ mg}\cdot\text{L}^{-1}$, $\text{pH} = 11$, $T = 70^\circ\text{C}$ and electrolysis time = 8 h.

Keywords: Electrochemical degradation; hydroxyl radicals; organic pollutants; lead dioxide; wastewater

Introduction

Dimethoate (O,O-dimethyl S-[2-(methylamino)-2-oxoethyl] phosphorodithioate) (Fig. 1) is a widely used organophosphorus (OPs) insecticide and acaricide applied to kill houseflies, as well as a wide range of insects and mites on a variety of fruit, vegetable, field and forestry crops. Like all OPs, dimethoate acts by interfering with the activities of cholinesterase, an enzyme essential for the proper functioning of the nervous system of insects and humans. A study showed that toxic effects and cholinesterase inhibition were observed in adult humans who ingested 30 mg a day or a higher dosage of dimethoate for 57 days (Hayes and Laws, 1990). Dimethoate is very toxic to birds, bees, fish and aquatic invertebrates (Cheminova, 1991). Dimethoate is degraded in the environment to another, more toxic, pesticide, omethoate; the proportion of omethoate in the total residue reaches about 50% after 5 weeks (FAO/WHO, 1985). Powerful degradation methods are required to destroy these pollutants in order to avoid their dangerous accumulation in the aquatic environment.

Among the different technologies for wastewater treatment, biological oxidation is frequently used. However, if wastewater contains highly toxic compounds, biological treatment may be useless. For this reason, there has been an increasing interest in the use of new methods such as electrochemical oxidation. It has been proven to be a promising and attractive technique for the effective oxidation of wastewater containing organic compounds (Chen, 2004; Fernandes et al., 2004; Sanroman et al., 2004). Many studies have demonstrated that the complete mineralisation of organics can be obtained with high efficiency by direct electro-oxidation using only high oxygen overvoltage anodes such as SnO_2 (Belhadj-Tahar and Savall, 1998;

Comninellis and Pulgarin, 1993; Polcaro et al., 1999 Stucki et al., 1991), PbO_2 (Andrade et al., 2007; Feng et al., 1995; Feng and Li, 2003; Martinez-Huitte et al., 2004a; Panizza and Cerisola, 2008; Quiroz et al., 2005; Samet et al., 2010a) and boron-doped diamond (BDD) anodes (Bechtold et al., 2006; Iniesta et al., 2001; Panizza and Cerisola, 2005; Samet et al., 2010b; Weiss et al., 2008). For example, some studies reported that the current efficiencies obtained with Si/BDD in oxidising 4-chlorophenol (Gherardini et al., 2001), 2-naphthol (Panizza and Cerisola, 2004), chloranilic acid (Martinez-Huitte et al., 2004a), and chlorpyrifos pesticide (Samet et al., 2010b) were higher than those obtained with PbO_2 anodes. In contrast, Martinez-Huitte et al. (2004b) showed that the oxidation of oxalic acid was faster at Ti/PbO_2 than at BDD, because the interaction of the oxalic acid with the PbO_2 surface was particularly strong, and its anodic oxidation was limited only by mass transfer at higher current densities and lower substrate concentrations. Recently, Flox et al. (2009) have demonstrated that *m*-cresol is more rapidly removed with PbO_2 than with BDD.

Both PbO_2 and BDD anodes exhibit good chemical and electrochemical stabilities, long lifetimes, and wide potential windows for water discharge. However, in the case of BDD, the financial expense can represent a serious drawback for industrial-scale wastewater treatment. In contrast, PbO_2 anodes are mainly used in the electrolytic production of perchlorates and chlorates (Kuhn and Wright, 1971). These anodes are electrochemically deposited on the metal surface. Titanium, niobium, tantalum and zirconium are widely used as substrate materials for their well-known properties, such as resistance to corrosion and chemical attack by acids, alkalis or salt solutions, and high mechanical strength.

Various innovative technologies have been proposed for treatment of wastewaters containing dimethoate. These include the use of photocatalytic oxidation, using TiO_2 as catalyst (Chen et al., 2007; Evgenidou et al., 2005), photo-Fenton process (Nikolaki et al., 2005), thermal decomposition (Andreozzi et al., 1999), ozonation (Liu et al., 2008), and microwave

* To whom all correspondence should be addressed.

✉ +21698660766; fax: +21674274437;

e-mail: youssef.samet@fss.rnu.tn

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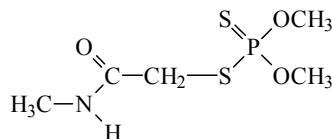


Figure 1
Chemical structure of dimethoate

irradiation of dimethoate in aqueous solutions (Zhang et al., 2007). Only one paper related to electrochemical oxidation, using Ti/Pt anodes, was found in the literature (Vlyssides et al., 2004). For this reason, we have opted to study the electrochemical degradation kinetics of dimethoate in aqueous solutions. The process was studied under galvanostatic polarisation mode in acidic media using Nb/PbO₂ anodes. The degradation rate of the studied pesticide and the limits of such a technique were investigated according to experimental parameters.

Experimental

Preparation of Nb/PbO₂ electrodes

Niobium surface treatment

Pre-treatments of the niobium substrate (rectangular plates 70 mm × 10 mm × 1 mm) were carried out before anodisation to ensure good adhesion of the lead dioxide film. Niobium was first roughened to increase the adhesion of PbO₂ deposit by subjecting its surface to mechanical abrasion using silica grains with an average diameter of 0.3 mm projected under 500 kPa pressure. It was then cleaned to remove sand particles or any other particles lodged on the metal surface. This process was carried out by degreasing with acetone, because of its ease of application and great penetrating power. The niobium was then ultrasonically rinsed in double-distilled water for 10 min. Achieving a uniform and well-adhered deposit requires a smooth surface with no oxides (formed spontaneously on contact with oxygen in the air) or scales. To ensure this, the niobium substrate was soaked for 30 s in hydrofluoric acid (40% weight) at room temperature, after which it was well-rinsed with double-distilled water. This process results in an average mass loss of niobium of about 0.156 mg·cm⁻².

Electrochemical deposition of PbO₂

The lead dioxide was deposited galvanostatically on the pre-treated niobium substrate by electrochemical anodisation of an aqueous Pb(NO₃)₂ solution (1 mol·l⁻¹) placed in a single-compartment Pyrex glass cell ($V=200\text{ cm}^3$) thermoregulated at 65°C. The cathode was a cylindrical platinum grid ($\phi=4\text{ cm}$, $L=6\text{ cm}$). The electrodeposition of PbO₂ film was carried out at an apparent current density of 10 mA·cm⁻² for 0.5 h, then at 20 mA·cm⁻² for the same period, and finally at 50 mA·cm⁻² for 1 h. The average mass of PbO₂ was 0.22 g·cm⁻². The deposit obtained was a grey porous material with strong adherence.

Chemicals

Dimethoate solutions were prepared from an emulsifiable concentrate (Biomat 40 EC, from Arysta Lifescience) containing 400 g·l⁻¹ dimethoate. All of the previous solutions were freshly prepared using double-distilled water and kept in the dark to avoid any photochemical reactions. Sulphuric acid and sodium hydroxide were of analytical grade and were purchased from Merck.

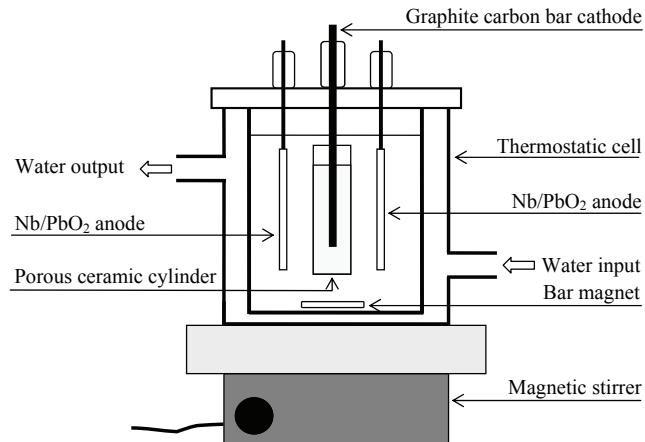


Figure 2
Electrolytic cell

Electrolysis

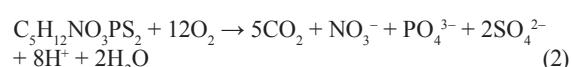
Galvanostatic electrolysis of dimethoate aqueous solutions (150 cm³) was carried out in a 2-compartment thermostatic cell (Fig. 2). The cathode was a graphite carbon-PTFE bar ($\phi=5\text{ mm}$; $L=60\text{ mm}$) placed in a porous ceramic cylinder (Norton, RA 84) containing 1 mol·l⁻¹ sulphuric acid solution. The anode is made up of 2 Nb/PbO₂ plates arranged symmetrically around the cathode; the surface of each plate not facing the cathode was masked with a protective film (transparent polyethylene type, Scotch TM 480, 3M). The geometric working surface of each plate was 5 cm². Dimethoate solutions were electrolysed in galvanostatic mode using a DC power supply (model ABTP 530 Française d'Instrumentation, France). The range of applied current density was 10 to 50 mA·cm⁻². The pH of the solution was adjusted, before and over the course of the electrolysis, by adding either concentrated sulphuric acid or sodium hydroxide solutions.

Analysis

Measurement of chemical oxygen demand (COD) during the process permitted evaluation of the kinetics of organic matter decay and mineralisation efficiency. Several authors used only COD measurements to follow organic matter decay during oxidation of organic compounds (Hmani et al., 2009; Panizza and Cerisola, 2008). In electrolysis using anodes with high oxygen evolution overpotential, such as PbO₂, the organic pollutants are mainly mineralised on the anode surface by the adsorbed hydroxyl radicals (HO[·]) electro-generated by water discharge (Eq. (1)).



The stoichiometry of dimethoate combustion indicates that 12 mol of O₂ are needed for the complete oxidation of dimethoate (Eq. (2)).



According to this reaction (Eq. (2)) the concentration of dimethoate (mol·m⁻³) in the electrolyte can be related to the COD (mol O₂·m⁻³) by the following relation:

$$[\text{C}_5\text{H}_{12}\text{NO}_3\text{PS}_2] = \frac{\text{COD}}{12} \quad (3)$$

When we compared the initial concentration of dimethoate in the solution prepared from the emulsifiable concentrate (Biomat 40 EC containing 400 g·l⁻¹ dimethoate) to that calculated from COD₀ (Eq. (3)), we found a difference of about 7%. In this study, during electrolysis, we used COD as an indicator for the presence of dimethoate and its by-products.

COD was determined by the dichromate method. The appropriate amount of sample was introduced into a prepared digestion solution (0–1 500 mg·l⁻¹) containing potassium dichromate, sulphuric acid and mercuric sulphate, and the mixture was then incubated for 2 h at 150°C in a COD reactor (WTW CR 2200 thermoreaktor Germany). COD concentration was measured colorimetrically using a DR/2010 spectrophotometer (Hach Company, USA).

The instantaneous current efficiency (ICE) for the anodic oxidation of the dimethoate was calculated from the values of the COD using the following relationship (Panizza and Cerisola, 2008):

$$\text{ICE}(\%) = FV \frac{\text{COD}_t - \text{COD}_{t+\Delta t}}{8I\Delta t} \times 100 \quad (4)$$

where:

F is the Faraday constant (96 485 C·mol⁻¹)

V is the volume of the solution (l)

COD_{*t*} and COD_{*t+Δt*} are the chemical oxygen demands

(g O₂·l⁻¹) at times *t* (s) and *t+Δt*, respectively

I is the applied current (A)

8 is the oxygen equivalent mass (g·mol⁻¹).

Results and discussion

Effect of the apparent applied current density

Figure 3 shows the trend of the COD/COD₀ ratio and ICE with time, for the anodic oxidation of dimethoate solution (COD₀ = 320 mg·l⁻¹) on the PbO₂ electrode, at different apparent current densities (10, 30 and 50 mA·cm⁻²). It is clear that COD/COD₀ ratio decreased almost exponentially with time and that the COD removal rate increased with increasing current. However, under these experimental conditions, the complete degradation of the organic matter was not reached. After 8 h of electrolysis, COD per cent removal increased from 54 to 81% when the apparent applied current density increased from 10 to 50 mA·cm⁻².

According to the literature (Belhadj-Tahar and Savall, 1998; Cominellis and Pulgarin, 1991; Samet et al., 2006; Torres et al., 2003), organic pollutants are directly destroyed by reaction with hydroxyl radicals formed at the anode surface from water oxidation (Eq. (1)). It is assumed that the hydroxyl radical concentration is constant during electrolysis; the COD removal rate *r* can be given by the following equation:

$$r = -\frac{d\text{COD}}{dt} = k [\text{HO}^{\cdot}]^{\alpha} \text{COD}^{\beta}(t) = k_{\text{app}} \text{COD}^{\beta}(t) \quad (5)$$

where:

α and *β* are the reaction orders related to the hydroxyl radicals and COD, respectively

k is the real rate constant

k_{app} is the apparent rate constant for COD removal.

If we supposed that COD removal kinetics followed a pseudo

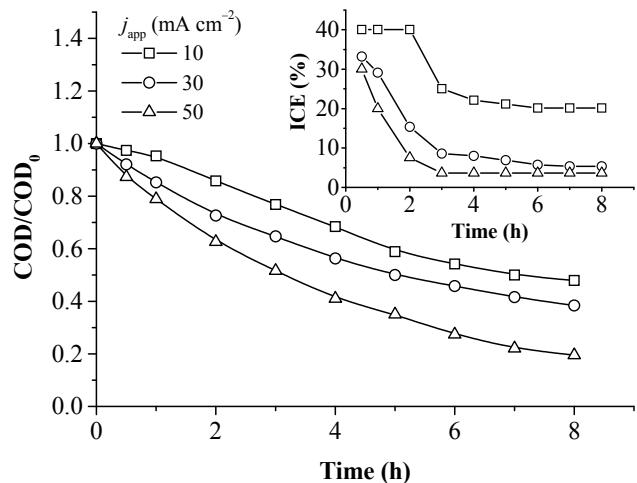


Figure 3
Influence of the apparent applied current density on the trend in COD/COD₀ ratio and ICE (inset) with time during the anodic oxidation of dimethoate (COD₀ = 320 mg·l⁻¹) at different apparent applied current densities. pH = 3 and T = 50°C.

first-order model ($\beta = 1$), the integration of Eq.(5) subject to the initial condition COD(*t*) = COD(0) at *t* = 0 leads to the following equation:

$$\ln \frac{\text{COD}_0}{\text{COD}} = k_{\text{app}} t = \frac{A}{V} k_m t \quad (6)$$

where:

k_m is the overall mass transport coefficient

A is the electrode surface (m²) and *V* is the volume of the solution (m³).

Figure 4 shows that the results are in agreement with Eq. (6), as indicated by the high value of the regression coefficient ($R^2 > 0.98$).

Using *k_m* values, the limiting current density (*j_{lim}⁰*) can be calculated with the initial COD value by the following equation (Panizza et al., 2001):

$$j_{\text{lim}}^0 = 4 F k_m \text{COD}_0 \quad (7)$$

Two kinetics regimes can be defined depending on the value of the applied current density (*j_{app}*):

- If *j_{app}* is lower than *j_{lim}⁰*, the electrolysis is under charge-transfer control; the instantaneous current efficiency is then 100%.
- If *j_{app}* is higher than *j_{lim}⁰*, the electrolysis is under mass-transport control; secondary reactions (such as oxygen evolution) occur.

Table 1 shows that *j_{lim}⁰* values were lower than *j_{app}* values; thus the electrolysis was under mass-transport control. This is in agreement with the trend for ICE calculated from Eq. (4), as shown in the inset of Fig. 3. ICE was always below 100% and decreased when *j_{app}* increased. ICE decreased significantly when *j_{app}* ranged from 10 to 30 mA·cm⁻² since α (*j_{app}/j_{lim}⁰*) was amplified approximately twice (α increased from 6.75 to 12.29, as shown in Table 1). However, there was no significant change in ICE values between 30 and 50 mA·cm⁻² current densities (inset of Fig. 3) due to the low variation of α within this range.

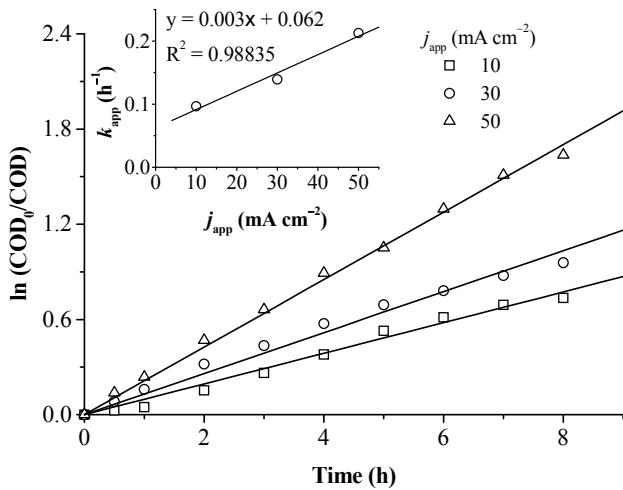


Figure 4

Fitting of the experimental data (for different j_{app}) to a first-order reaction kinetic model. The inset panel shows k_{app} evolution at different j_{app} . Dimethoate ($COD_0 = 320 \text{ mg} \cdot \text{l}^{-1}$); pH = 3 and $T = 50^\circ\text{C}$.

Table 1 Apparent rate constant (k_{app}), overall mass transport coefficient (k_m) and initial limiting current density (j_{lim}^0) as a function of the apparent applied current densities			
j_{app} (mA·cm ⁻²)	10	30	50
$k_{app} \times 10^2$ (h ⁻¹)	9.20	15.20	21.20
$k_m \times 10^6$ (m·s ⁻¹)	3.83	6.33	8.83
j_{lim}^0 (mA·cm ⁻²)	1.48	2.44	3.41
$\alpha = j_{app} / j_{lim}^0$	6.75	12.29	14.67

Effect of the initial concentration of dimethoate

Figure 5 shows the trend of COD removal with time during the anodic oxidation of dimethoate at different initial concentrations (COD_0 : 100, 320, and 550 $\text{mg} \cdot \text{l}^{-1}$) using an apparent current density of 50 $\text{mA} \cdot \text{cm}^{-2}$, at pH 3 and 50°C . The results indicated that COD per cent removal decreased with an increase in initial COD. After 8 h of electrolysis, COD per cent removal decreased from 85 to 65% when initial COD increased from 100 to 550 $\text{mg} \cdot \text{l}^{-1}$.

Considering a pseudo first-order reaction for COD removal, k_{app} values calculated from the straight lines decreased progressively from $27.72 \cdot 10^{-2}$ to $14.31 \cdot 10^{-2} \text{ h}^{-1}$ when the initial COD ranged from 100 to 550 $\text{mg} \cdot \text{l}^{-1}$, as shown in Table 2. However, overall mass transport rate ($k_m \times COD_0$) increased with increased concentrations of dimethoate. Consequently, the COD removal rate was faster when the initial COD was higher. This process can be interpreted in terms of an increase in the overall mass transport rate to the anode surface. In fact, when COD_0 increased, the rate of formation and detachment of O_2 bubbles (Eq. (8)) decreased. Thus the mass transport rate can be significantly increased.



Effect of temperature

The electrolysis of dimethoate solutions ($COD_0 = 320 \text{ mg} \cdot \text{l}^{-1}$) at the PbO_2 anode, at an apparent applied current density of

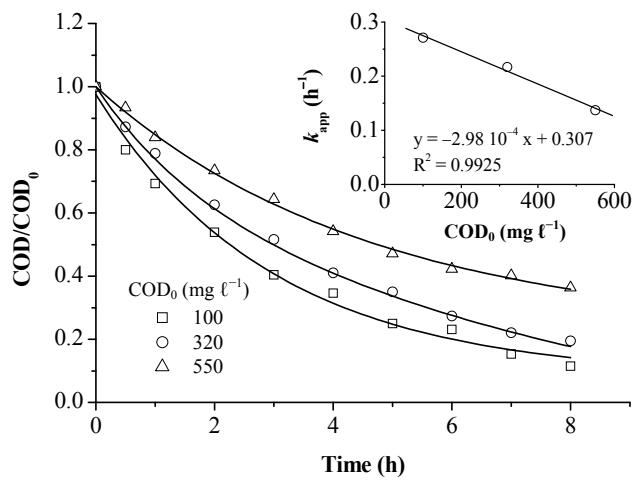


Figure 5

Influence of the initial concentration of dimethoate (given by COD_0) on the trend in COD/COD_0 ratio during electrolysis. The inset panel shows k_{app} evolution at different COD_0 . pH = 3; $T = 50^\circ\text{C}$ and $j_{app} = 50 \text{ mA} \cdot \text{cm}^{-2}$.

Table 2 Apparent rate constants (k_{app}), overall mass transport coefficient (k_m) and initial overall mass transport rate ($k_m \times COD_0$) as a function of the initial COD. COD_1 and COD_8 are COD values after 1 and 8 h electrolysis time, respectively.			
$COD_0(\text{mg} \cdot \text{l}^{-1})$	100	320	550
$COD_1(\text{mg} \cdot \text{l}^{-1})$	56	252	423
$k_{app} \times 10^2$ (h ⁻¹)	27.72	21.20	14.31
$k_m \times 10^6$ (m·s ⁻¹)	11.55	8.83	5.96
$k_m \times COD_0 \times 10^2$ (mg·m ⁻² ·s ⁻¹)	115	282	327
$k_m \times COD_1 \times 10^2$ (mg·m ⁻² ·s ⁻¹)	64	222	252
COD_8 removal %	86	80	65

50 $\text{mA} \cdot \text{cm}^{-2}$, was carried out at different temperatures, in the range of 30–70°C. Figure 6 shows that an increase in temperature had a positive effect on the COD removal rate. The amount of organic matter decreased more quickly at 70°C than at 30°C. After 8 h of electrolysis, COD per cent removal increased from 22% to 93% when the temperature ranged from 30 to 70°C. This result is probably due to the increase in the indirect oxidation reaction of organics by the electro-generated oxidising agent $S_2O_8^{2-}$ from the supporting electrolyte H_2SO_4 (Eq. (9)).



Several authors have shown that $S_2O_8^{2-}$ formed at the PbO_2 anode surface participates in the oxidation of the organic matter in proximity to the electrode surface and/or in the bulk of the solution (Amadelli et al., 2002; Aquino et al., 2010; Flox et al., 2009; Liu and Liu, 2008). This results in an increase in COD removal rate with temperature. These results are in agreement with those obtained by other authors (Canizares et al., 2005; Michaud et al., 2000). Moreover, the decrease in the viscosity of the medium with increase in temperature enhances the diffusion rate of organic matter to the anode surface.

Consequently, an increase in temperature plays a role in an increase in COD removal rate. The same explanation was suggested by Rodriguez et al. (2009).

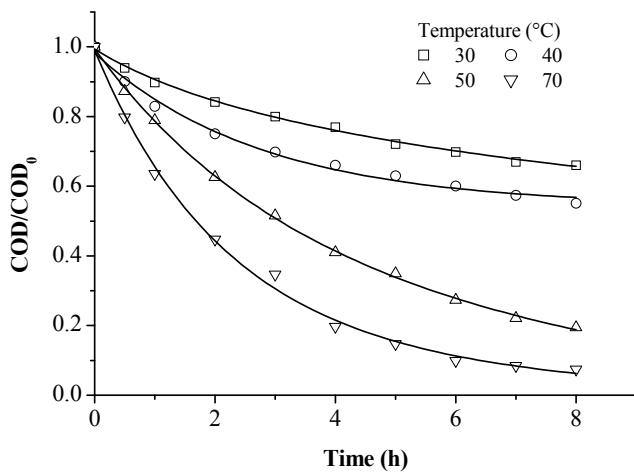


Figure 6

Influence of temperature on the trend in COD/COD_0 ratio with time during the anodic oxidation of dimethoate ($\text{COD}_0 = 320 \text{ mg}\cdot\text{L}^{-1}$) at different temperatures. $\text{pH} = 3$ and $j_{\text{app}} = 50 \text{ mA}\cdot\text{cm}^{-2}$.

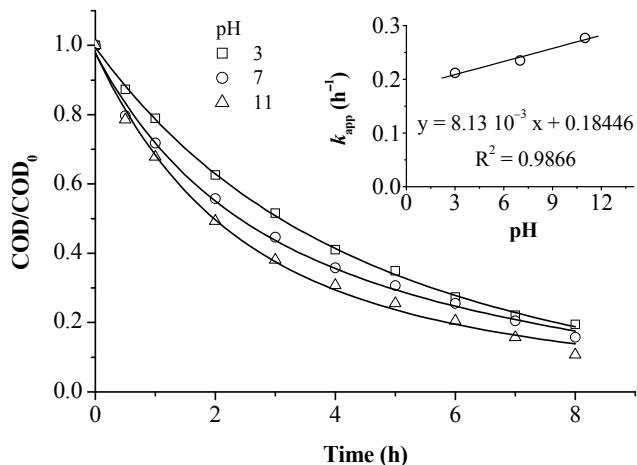


Figure 8

Influence of pH on the trend in COD/COD_0 ratio with time during the anodic oxidation of dimethoate ($\text{COD}_0 = 320 \text{ mg}\cdot\text{L}^{-1}$) at different pH values. The inset panel shows k_{app} evolution at different pH. $j_{\text{app}} = 50 \text{ mA}\cdot\text{cm}^{-2}$ and $T = 50^\circ\text{C}$.

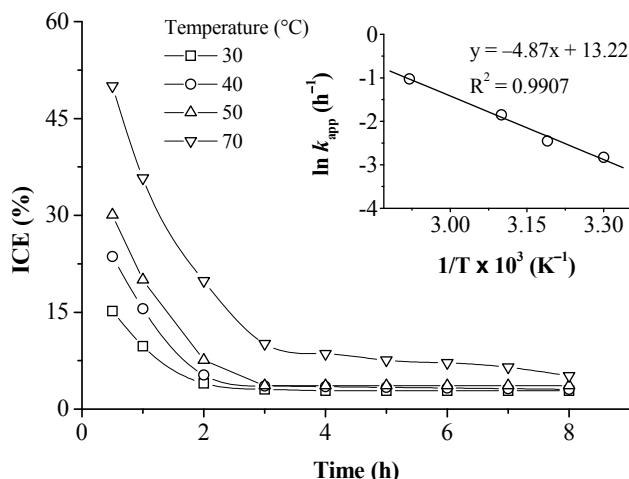


Figure 7

ICE curves for a range of temperatures during electrolysis of dimethoate ($\text{COD}_0 = 320 \text{ mg}\cdot\text{L}^{-1}$). The inset panel shows the Arrhenius-type plot of k_{app} . $\text{pH} = 3$ and $j_{\text{app}} = 50 \text{ mA}\cdot\text{cm}^{-2}$.

From Fig. 7, we can note that ICE values were more significant at a high temperature (70°C). Thus, the degradation reaction of the organic matter predominated. The Arrhenius expression, showing the relationship between the reaction temperature and k_{app} is expressed as follows:

$$k_{\text{app}} = A \exp\left(-\frac{E_{\text{app}}}{R T}\right) \quad (10)$$

where:

- A is the pre-exponential (or frequency) factor
- E_{app} is the apparent global activation energy ($\text{J}\cdot\text{mol}^{-1}$)
- R is the ideal gas constant ($8.314 \text{ J}\cdot\text{mol}^{-1}\cdot\text{K}^{-1}$)
- T is the absolute temperature (K).

Due to the narrow temperature range employed in this study (30 – 70°C), variations of the pre-exponential factors and the apparent activation energy of the empirical Arrhenius expressions of the COD removal may be ignored. The variation of

$\ln k_{\text{app}}$ versus $1/T$ is plotted in the inset of Fig. 7. A good linear relationships exists between the plot of $\ln k_{\text{app}}$ and $1/T$. Based on the slope ($-E_{\text{app}}/R$) and intercepts ($\ln A$) of the plot in Fig. 7, E_{app} and A in Arrhenius form (Eq. (10)) were determined as $35.58 \text{ kJ}\cdot\text{mol}^{-1}$ and 152.78 s^{-1} , respectively. E_{app} for a diffusion-controlled homogeneous reaction is typically less than $40 \text{ kJ}\cdot\text{mol}^{-1}$ (Belhadj-Tahar and Savall, 1998). The experimental result was considerably lower than this value; it is therefore likely that the limiting step of dimethoate oxidation is of a diffusional nature.

Effect of pH

Many studies have reported on the effects of solution pH in anodic oxidation of organics on PbO_2 anodes, 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 packed-bed anode increased from 3% to 13% as pH increased from 2 to 11. In contrast, in a previous study (Samet et al., 2006) we demonstrated that the oxidation kinetics for 4-chloroguaiaicol removal using PbO_2 anodes were faster at pH 2.0 than at pH 6.0. Recently, 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. The same result was observed by Chen and Chen (2006) when they studied the oxidation of orange II at a BDD anode. These discrepancies can be explained by the differences in the chemical properties of the organic compounds tested. To study the influence of pH in the oxidation of dimethoate, electrolyses were carried out at 3 initial pH values in the range of 3–11 (Fig. 8). It was found that, within the pH range studied, COD removal was slightly affected by solution pH. After 8 h, COD per cent removal increased from 80 to 90% with an increase in pH from 3 to 11. A pH of 11 appeared to be the optimum value. The inset of Fig. 8 shows a slight increase in k_{app} with increasing solution pH.

Conclusions

The electrocatalytic activity of PbO_2 anode for the electrochemical oxidation of an aqueous solution of dimethoate was

investigated under different experimental conditions. The results of this study reveal the following traits:

- 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, in the potential region of water oxidation.
- Organics degradation was a diffusion-controlled process with a higher rate of COD removal obtained at high apparent applied current density ($50 \text{ mA} \cdot \text{cm}^{-2}$).
- An increase in temperature significantly improved the COD removal rate. For $\text{COD}_0 = 320 \text{ mg} \cdot \text{l}^{-1}$, $j_{\text{app}} = 50 \text{ mA} \cdot \text{cm}^{-2}$, $\text{pH} = 3$ and $T = 70^\circ\text{C}$, 90% of COD removal was achieved after 8 h.
- Dimethoate degradation was favoured when operating at low initial COD.
- Solution pH had only a slight positive effect on COD removal.

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