

Modelling the change in the oxidation coefficient during the aerobic degradation of phenol by acclimated activated sludge

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ABSTRACT

In this work the aerobic degradation of phenol by acclimated activated sludge was studied. Results demonstrate that while the phenol removal rate by acclimated activated sludge follows the Monod model, the oxygen uptake rate obeys a Haldane-type equation. The phenol oxidation coefficient obtained at different initial phenol concentrations ranged from 1.9 to 2.6 mol O₂ · mol⁻¹ phenol. A mathematical model based on a simplified version of the catalytic mechanism of the enzyme phenol 2-monoxygenase was developed to predict transient phenol concentrations and oxygen requirements by phenol-acclimated activated sludge in batch reactors under different initial phenol concentrations. The proposed model not only adequately represents the experimental results of the present paper, but also those reported by other authors. Particular cases of the proposed model are discussed.

Keywords: oxygen uptake rate; phenol; biodegradation; biokinetic model; respirometry

INTRODUCTION

Phenolic compounds are considered to be a major group of hazardous environmental pollutants. Several industries, such as petroleum processing plants, oil refineries, coke oven and pharmaceuticals, generate large amounts of phenolic wastewaters (Lepik and Tenno, 2011; Pramparo et al., 2012). Phenol is used as an intermediate in the production of phenolic resins, which are used in the plywood, adhesive, construction, automotive and appliance industries, in the production of synthetic fibres and for epoxy resin precursors. Besides, phenolic compounds are naturally present in some foodstuffs, in human and animal wastes, and in decomposing organic material, and are produced from the metabolism of proteins (Lin and Juang, 2009).

Phenols can be removed from industrial effluents by several physicochemical methods. Separation technologies include distillation, liquid–liquid extraction, adsorption over activated carbons and polymeric and inorganic adsorbents, membrane pervaporation and membrane–solvent extraction. Destruction technologies such as non-catalytic, supercritical and catalytic wet air oxidation, ozonation, non-catalytic, catalytic and enzymatic peroxide wet oxidation, electrochemical and photocatalytic oxidation have also been studied. As a general rule, all these treatments are usually complex and very expensive; for these reasons, biological methods are preferred (Busca et al., 2008).

Many aerobic bacteria and mixed cultures are capable of degrading aromatic compounds as the sole carbon and energy source (El-Naas et al., 2009; Banerjee and Ghoshal, 2010). Batch and semi-continuous systems with suspended or immobilised

biomass were used to study phenol degradation kinetics (Orupold et al., 2001; Nuhoglu and Yalcin, 2005; Tziotziou et al., 2005). The most used model to describe the dependence of biomass growth, phenol and oxygen consumption rates on a self-inhibitory substrate, such as phenol, is the Haldane equation with a constant yield (Christen et al., 2012; Pramparo et al., 2012). Although the Haldane model can be derived from an enzymatic mechanism, not all the enzymes involved in phenol metabolism can be represented by this mechanism. In particular, the first step in the aerobic biodegradation of phenol comprises the oxidation of phenol to catechol by the enzyme phenol 2-monoxygenase (PH2MO). The catalytic mechanism of PH2MO is complex, involving FAD and 3 substrates (molecular oxygen, phenol and NADPH) (Enroth et al., 1998).

Due to the lack of fit of the traditional approach (e.g., the Haldane equation with a constant yield), several authors (Feitkenhauer et al., 2001; Nuhoglu and Yalcin, 2005; Ben-Youssef and Vázquez-Rodríguez, 2011) have proposed different modifications to the traditional model. However, those models were developed on an empirical base, i.e. the study of phenol degradation and biomass growth kinetics and then the fitting of an appropriate mathematical model to the obtained results. Additionally, most studies concerning the aerobic degradation of phenol have been performed using pure cultures, low concentration ranges of phenol, or even a single initial concentration of phenol. In this sense, further investigation is necessary to understand phenol biodegradation under more realistic conditions, especially in biological wastewater treatment systems where the concentration of phenol can vary over a wide range (Alcocer et al., 2007; Banerjee and Ghoshal, 2010; El-Naas et al., 2010).

Knowledge of microbial substrate consumption kinetics is important for accurate prediction of the quality of the treatment process effluent. One key feature is the relationship between phenol concentration and the rates of substrate and oxygen consumption (Kumar et al., 2005; Nuhoglu and Yalcin,

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2005). While the substrate consumption rate determines the size of bioreactors, knowledge of the oxygen uptake rate associated with phenol degradation is crucial to an appropriate design of aeration devices. Taking into account that, in a typical activated sludge wastewater treatment plant, aeration costs represent approximately 45% of the total operative cost (e.g. energy for aeration, pumping, and sludge treatment) (Vanrolleghem and Gillot, 2002), accurate kinetic models help engineers to optimise operational conditions in order to meet discharge requirements, minimising operational costs. For these reasons, in this work a mathematical model based on a simplified version of the catalytic mechanism of PH2MO was developed to predict transient phenol concentrations and oxygen requirements by phenol-acclimated activated sludge in batch reactors under different initial phenol concentrations.

EXPERIMENTAL

Chemicals and reagents

Phenol (loose crystals, > 99%) was obtained from Sigma (St. Lois, MO, USA). All inorganic salts were commercial products of reagent grade from Anedra (San Fernando, Argentina). Dehydrated cheese whey was from Food S.A. (Villa Maipú, Argentina).

Acclimation of activated sludge to phenol

Activated sludge used in this study was cultured in a laboratory-scale (2.5 l) cylindrical semicontinuous fill and draw reactor. Aeration was provided at the bottom of the reactor through an air-stone using 2 air pumps at 2 l·min⁻¹; dissolved oxygen (DO) concentration was maintained above 4 mg·l⁻¹. Phenol-acclimated activated sludge was obtained as follows: Initially, the reactor was fed with a model wastewater of the dairy industry (Lobo et al., 2013), dehydrated cheese whey 1 000 mg, (NH₄)₂SO₄ 940 mg, K₂HPO₄ 500 mg and KH₂PO₄ 250 mg. All components were diluted in 1 l of tap water. The hydraulic retention time was 80 h and a solids retention time of 30 days was maintained by direct wastage. After 70 days of operation, the model wastewater was replaced by a culture medium with phenol as the sole carbon-limiting source (Nuhoglu and Yalcin, 2005): (NH₄)₂SO₄ 226 mg·l⁻¹, K₂HPO₄ 500 mg·l⁻¹, KH₂PO₄ 250 mg·l⁻¹, MgSO₄·7H₂O 25.2 mg·l⁻¹, MnSO₄·H₂O 2.52 mg·l⁻¹, CaCl₂ 2 mg·l⁻¹, FeCl₃ 1.2 mg·l⁻¹, phenol 300 mg·l⁻¹. pH was adjusted to 7.0 ± 0.05 by adding a few drops of concentrated solutions of NaOH or HCl. During this phase the solids retention time was increased to 45 days in order to prevent biomass washout. The reactor was monitored periodically by measurements of total suspended solids (TSS), soluble chemical oxygen demand (sCOD) and soluble COD consumption rate (R_{COD}). Biomass was considered acclimated to phenol when the specific sCOD consumption rate (q_{COD} = R_{COD}/TSS) was approx. constant for at least 2 weeks.

During several fill and draw cycles, total phenol (TPh) and DO were also measured as a function of the cycle time. In these cases, before the addition of phenol, the oxygen mass transfer coefficient (k_La) was obtained using a non-steady state procedure (Ros, 1993). The oxygen uptake rate associated with phenol consumption (R_{O₂}) was calculated from the DO mass balance in the reactor:

$$R_{O_2} = k_L a (C_E - C) - \frac{dC}{dt} \quad (1)$$

where:

C_E is the DO concentration in equilibrium with respiration in the absence of substrate

C is the instantaneous DO concentration

The amount of oxygen consumed (OC) during the phenol oxidation was calculated as follows:

$$OC = \int_0^t R_{O_2} dt \quad (2)$$

The overall oxidation coefficient (Y_{O/S}), which represents the amount of oxygen consumed per unit of substrate (phenol) oxidised, was calculated as the ratio OC_m/S₀, where OC_m is the maximum amount of oxygen consumed and S₀ is the initial phenol concentration (mg·l⁻¹).

Effect of the initial phenol concentration on the biodegradation kinetics

The effect of initial phenol concentration on phenol degradation kinetics by phenol-acclimated activated sludge was assessed using an open (flowing gas/static liquid) respirometer. Before performing the assays, activated sludge samples were washed 3 times with phosphate buffer (15 mM, pH = 7) and re-suspended to a final volume of 500 ml using the above-mentioned buffer. Then, 0.5 ml micronutrient solutions were added (Contreras et al., 2011). Washed activated sludge samples were poured into the respirometer, which consists of a 500-ml working volume reactor with temperature control (25 ± 0.5°C). The sample was stirred with a magnetic stir-bar and was aerated continuously by an air pump. Air was set to a stable flow rate (1.0 l·min⁻¹) using a high precision rotameter (Bruno Schilling model MB 60V, Argentina), and DO concentration as a function of time was recorded. Oxygen mass transfer coefficient of the respirometer (k_La) and R_{O₂} values were calculated as described in the previous section. In these experiments the biomass concentration was 0.7 g·l⁻¹ TSS and tested initial phenol concentration (S₀) ranged from 0.53 to 5.3 mM.

Analytical procedures

Total suspended solids (TSS) were used as a measure for the biomass concentration (X, g·l⁻¹ TSS). Duplicate biomass measurements were performed to reduce experimental errors; average and maximum relative errors for TSS were 4% and 13%, respectively. Dissolved oxygen (DO) concentration was measured using a polarographic DO probe YSI (model 5739). DO data were acquired by a personal computer interfaced to a DO monitor YSI (model 58). Soluble COD (sCOD) was determined as follows: 3 ml of culture samples were centrifuged for 5 min at 13 000 r·min⁻¹ (Eppendorf 5415C); then, COD of the supernatant was determined using a commercial test (Hach). Sample digestion (2 h, 150°C) was performed in a Hach COD Reactor 45600; a Hach DR 2000 photometer was used for the absorbance determination step. Total phenol (TPh) concentration of the supernatant was measured using the 4-aminoantipyrine (4-AAP) method (Vazquez-Rodriguez et al., 2006). This method uses 2 reagents, 4-AAP (20.8 mM of 4-AAP in 0.25 M NaHCO₃) and ferricyanide (83.4 mM of K₃Fe(CN)₆ in 0.25 M NaHCO₃), as colour-generating substrates when combined with phenolic compounds. Coloured complexes were measured at 510 nm in a Hach DR 2000 spectrophotometer. Calibration curves were performed periodically using phenol as the reference compound.

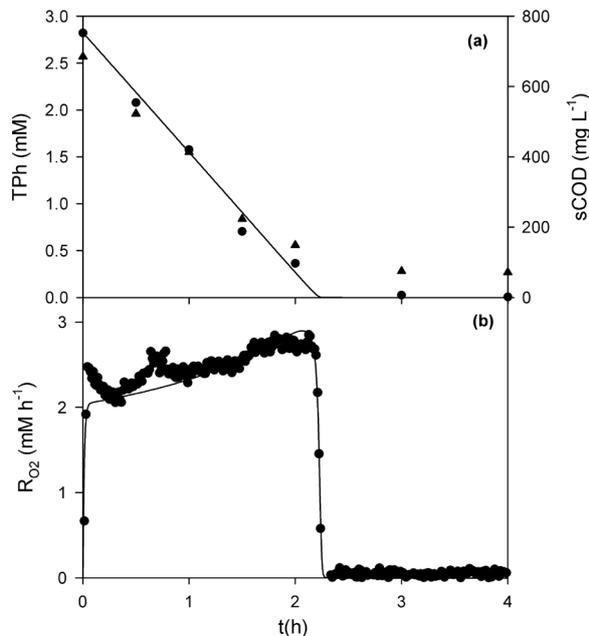


Figure 1

Example of (a) total phenols (TPh, ●), soluble COD (sCOD, ▲) and (b) oxygen uptake rate (R_{O_2} , ●) as a function of time during a fill and draw cycle. Solid lines indicate the results obtained by the proposed model (Eqs. (12) and (19)) using the coefficients shown in Table 1. In this example the biomass concentration was $1.8 \text{ g} \cdot \text{L}^{-1}$ TSS.

Estimation of the model coefficients and dynamic simulations

Coefficients of the mathematical model proposed in this work and the dynamic simulations were performed using the software package Gepasi 3. The routine LSODA (Livermore Solver of Ordinary Differential Equations) was selected to integrate the system of differential equations of the proposed model. Multistart Optimisation algorithm (with Levenberg-Marquardt local optimisation) was selected to fit the proposed model to the experimental data. In order to reduce fitting errors of the adjustable coefficients, initial concentrations were adjusted within $\pm 5\%$ of their measured values. This procedure takes into account the degree of uncertainty in the initial conditions due to analytical errors. To describe the biological start-up phenomenon, a first-order correction factor was included (Contreras et al., 2008).

RESULTS AND DISCUSSION

Phenol consumption and oxygen uptake by phenol-acclimated activated sludge

Figure 1a shows an example of total phenol (TPh) and soluble COD (sCOD) concentration profiles during a fill and draw cycle with phenol. Both TPh and sCOD exhibited a linear decrease up to phenol depletion at $t = 2.2$ h; within this interval, removal rates of TPh and sCOD were $1.26 \text{ mM} \cdot \text{h}^{-1}$ and $274 \text{ mg} \cdot \text{L}^{-1} \cdot \text{h}^{-1}$, respectively. Figure 1a shows that TPh removal is linear until its depletion, which corresponds to a constant removal rate of TPh. The oxygen uptake rate associated with phenol consumption (R_{O_2}) is shown in Fig. 1b. After a short start-up period, R_{O_2} increased from 2 to 3 $\text{mM} \cdot \text{h}^{-1}$ as a function of time; then, R_{O_2} decreased to values close to zero due to phenol depletion. Taking into account that during this experiment the biomass

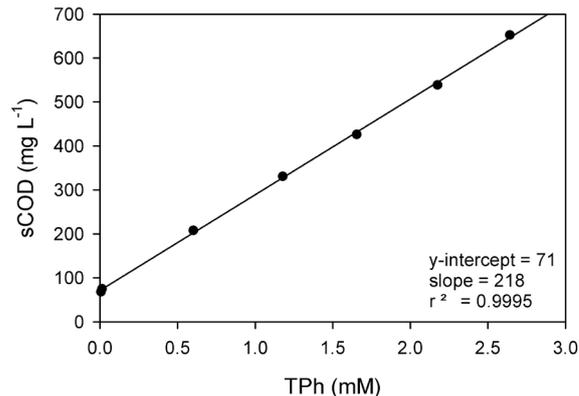


Figure 2

Soluble COD (sCOD) as a function of total phenol (TPh) corresponding to the example depicted in Fig. 1.

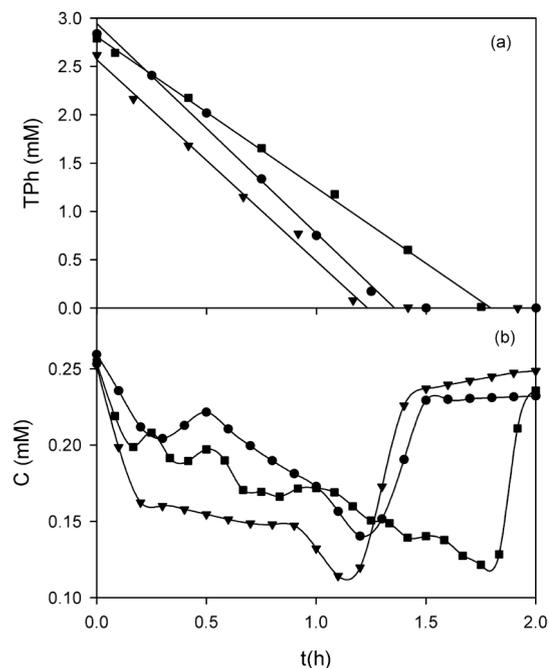


Figure 3

(a) Total phenol (TPh) and (b) dissolved oxygen (C) concentrations as a function of time during fill and draw corresponding to 126 (●), 130 (▼) and 160 (■) days of operation. Continuous lines in (a) indicate the linear regressions.

concentration (X) increased about 17% with respect to the initial value (from 1.8 to 2.1 $\text{g} \cdot \text{L}^{-1}$ TSS), the change of R_{O_2} values could not be attributed to the increment of biomass but to an intrinsic feature of the phenol biodegradation kinetics, such as the inhibition of R_{O_2} by phenol.

Figure 2 shows that the plot of sCOD as a function of TPh was a straight line with slope = $218 \text{ g COD} \cdot \text{mol}^{-1} \text{ TPh}$; because this slope value is close to the theoretical oxygen demand of phenol (e.g., $224 \text{ g COD} \cdot \text{mol}^{-1}$ phenol), it was concluded that no partial oxidation products of phenol were released during its degradation. When the phenol was depleted, the lowest sCOD concentration was about $71 \text{ mg} \cdot \text{L}^{-1}$; this background sCOD value was attributed to the presence of microbial soluble products (Xie et al., 2012).

Figure 3 shows several examples of total phenols (TPh) and dissolved oxygen (C) concentrations as a function of time

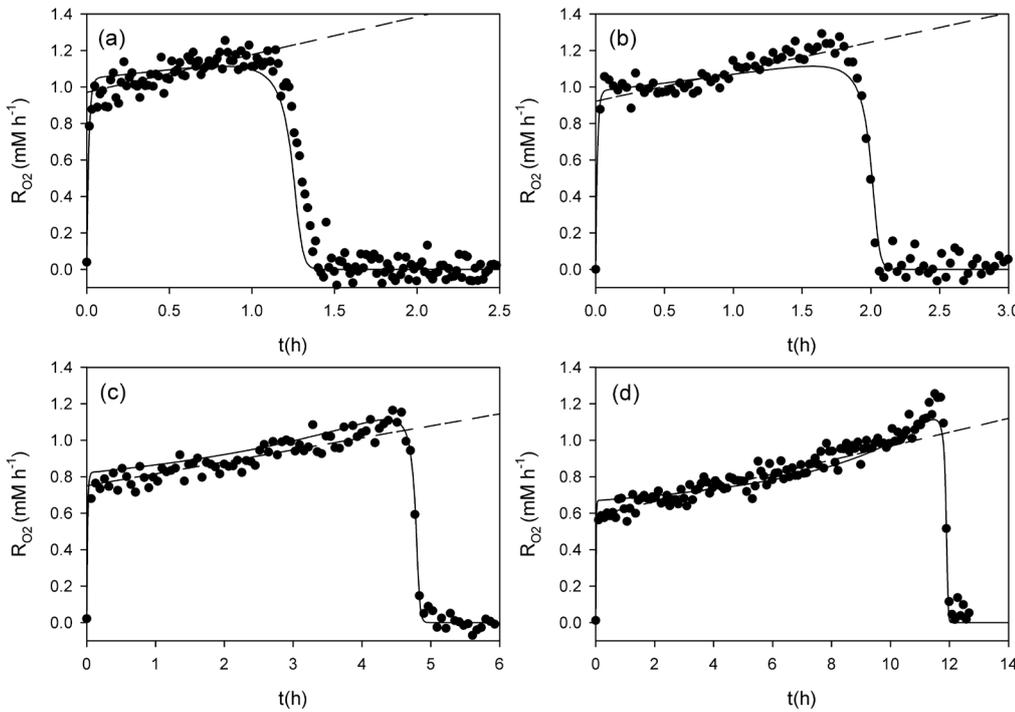


Figure 4
Oxygen uptake rates (R_{O_2}) as a function of time for different initial phenol concentrations (in mM): (a) 0.53, (b) 1.06, (c) 2.12, (d) 5.32. Continuous lines indicate the proposed model. Dashed lines indicate the regression line used to calculate the initial oxygen uptake rate (R_{O_20}). In these experiments the biomass concentration was $0.7 \text{ g} \cdot \ell^{-1}$ TSS.

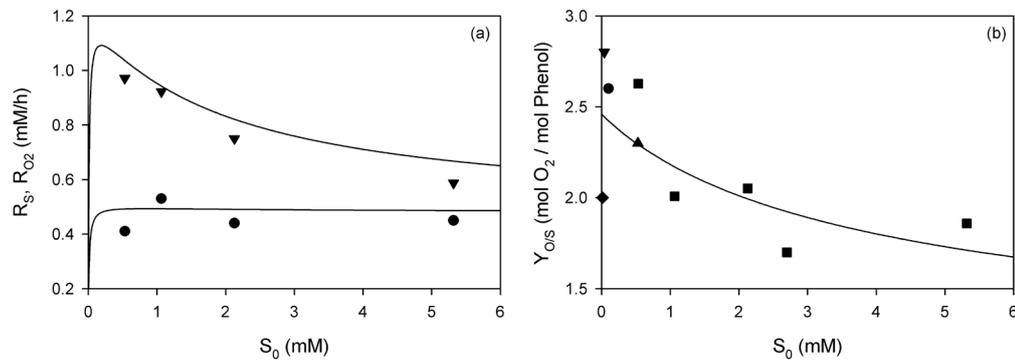


Figure 5
Effect of the initial phenol concentration (S_0) on (a) phenol (R_S , circles) and oxygen (R_{O_2} , triangles) consumption rates (a) and on (b) the overall oxidation coefficient ($Y_{O/S}$) measured in this work (squares) and by other authors: Chudoba et al. (1985) (●), Orupold et al. (2001) (▼), Contreras et al. (2008) (▲), Lepik and Tenno (2011) (◆). Solid lines indicate the results obtained by the proposed model using the coefficients shown in Table 1.

during a fill and draw cycle with phenol. In all cases, a linear decrease of *TPh* concentration was obtained (Fig. 3a). Taking into account that C values ranged between 0.10 and 0.25 mM (Fig. 3b), these results suggest that within the tested conditions the phenol consumption rate (R_S) was not affected by the dissolved oxygen concentration. Figure 3 also shows that the time at which phenol is depleted (t_F) is similar to the time at which C begins to increase; thus, the increase of C can be used as an indication of substrate depletion. Buitron et al. (2005) developed a control strategy to optimise the reaction time of a SBR based on the DO profile; according to those authors, the time at which the substrate is completely consumed (t_F) can be assessed by the increase in DO.

Effect of the initial phenol concentration on the biodegradation kinetics

Open respirometer experiments were performed in order to study the effect of initial phenol concentration on respiration rate and on oxygen consumed during phenol biooxidation. Figure 4 shows that for all the tested initial phenol concentrations ($S_0 = 0.53$ to 5.3 mM), after a start-up period R_{O_2} increased almost linearly as a function of time and then decreased to values close to zero due to substrate depletion. According to

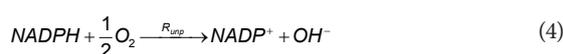
Fig. 3, the time at which the substrate is completely consumed (t_F) can be assessed by the increase in DO; based on these t_F values, average substrate consumption rates (R_{S-ave}) corresponding to different initial phenol concentrations were calculated as the ratio S_0/t_F . Figure 5a shows that R_{S-ave} values were constant as a function of the initial phenol concentration (S_0), suggesting Monod type kinetics for the removal rate of *TPh* (R_S). This result is in accordance with the linear decrease of *TPh* and *sCOD* depicted in Figs. 1a and 3a. Besides, from the extrapolation of R_{O_2} to time zero (Fig. 4), initial R_{O_2} values (R_{O_20}) corresponding to the different initial phenol concentrations can be obtained. Figure 5a shows that R_{O_20} decreased as S_0 increased, suggesting a Haldane dependence of R_{O_2} as a function of S . Note that if R_S follows the Monod equation and R_{O_2} obeys a Haldane-type dependence with respect to S , then the oxidation coefficient (e.g., the amount of oxygen consumed per unit of substrate oxidised) must be a function of S_0 , rather than a constant.

Based on the R_{O_2} profiles corresponding to different S_0 values (Fig. 4), maximum values for the oxygen consumed (OC_m) were calculated using Eq. (2). Figure 5b shows that the overall oxidation coefficient ($Y_{O/S}$) decreased as S_0 increased, confirming that the amount of oxygen consumed per unit of substrate oxidised was a function of S_0 . The range of $Y_{O/S}$ values corresponding to phenol obtained in the present work (1.9–2.6 mol

O₂·mol⁻¹ phenol) was similar to that reported by other authors. Orupold et al. (2001) studied the oxidation of several phenolic compounds by activated sludge; those authors reported a $Y_{O/S}$ value of 2.6 ± 0.5 mol O₂·mol⁻¹ phenol. Recently, Lepik and Tenno (2011) studied the aerobic degradation of phenol by activated sludge acclimated to a wastewater of an oil-shale industry; those authors reported a $Y_{O/S}$ value of 2.0 mol O₂·mol⁻¹ phenol. It should be noted that $Y_{O/S}$ values, calculated according to carbon and reduction degree balances applied to biomass growth with phenol as sole carbon source, range between 4.0 and 7.3 mol O₂·mol⁻¹ phenol (Feitkenhauer et al., 2001; Kumar et al., 2005). These differences suggest that respirometric $Y_{O/S}$ values reflect a partial oxidation of phenol due to the accumulation of metabolic intermediates. Wang and Loh (1999) studied phenol degradation by *Pseudomonas putida* ATCC 49451; those authors found that the accumulation of metabolic intermediates of phenol inhibited phenol oxidation, especially at high initial phenol concentrations. Similar results were reported by other authors with regard to the accumulation of metabolic intermediates during phenol degradation by activated sludge (Nuhoglu and Yalcin, 2005; Vazquez-Rodriguez et al., 2006, Ben-Youseff and Vazquez-Rodriguez, 2011).

Modelling phenol oxidation by acclimated activated sludge

The first step in the aerobic degradation pathway of phenol comprises the hydroxylation of phenol by the enzyme phenol 2-monooxygenase (PH2MO). In the present model it was assumed that the total PH2MO concentration (E_T) is proportional to the biomass concentration (X), being q_E , the proportionality constant. According to several authors, the catalytic mechanism of PH2MO involves 3 substrates: molecular oxygen, phenol and NADPH. The reaction scheme for PH2MO can be divided into reductive and oxidative half-reactions. During the reductive step, after the phenol binding, the oxidised form of the enzyme ($E_{ox}S$) is reduced by NADPH, NADP⁺ leaves the enzyme to produce an intermediary complex $E_{red}S$. In the oxidative half-reaction, molecular oxygen reacts with $E_{red}S$, phenol (S) is hydroxylated to catechol (SOH) and the oxidised form of the enzyme E_{ox} is regenerated (Enroth et al., 1998). Besides, Divari et al. (2003) demonstrate that during the catalytic cycle of PH2MO, the ratio of product formed to NADPH consumed is less than 1. Several authors (Enroth et al., 1998; Divari et al., 2003) have proposed that the mechanism of action of PH2MO must bifurcate into productive and unproductive paths, which involves the dissociation of $E_{red}S$ to form free reduced enzyme (E_{red}). The free reduced enzyme is then oxidised by oxygen to produce E_{ox} and H₂O₂. Because the decomposition of hydrogen peroxide returns one half of the oxygen consumed, the net result of the unproductive path is the consumption of 0.5 mol of oxygen. The overall reactions for the productive and unproductive paths are the following:

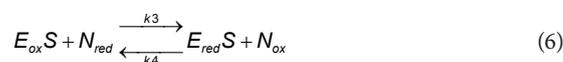


where:

R_s and R_{unp} are the rates of the productive and unproductive paths, respectively

$$d = \frac{k_2 k_6 k_p + k_{unp} k_1 (k_p + k_5) + N_T \{k_4 (k_2 k_6 + k_{unp} k_1) (1 - f_{red}) + k_3 [k_6 k_p + k_1 (k_{unp} + k_5) f_{red}]\}}{k_1 k_6 \{k_p + N_T [k_4 (1 - f_{red}) + k_3 f_{red}]\}} \quad (16)$$

Results shown in Fig. 3 suggest that within the experimental conditions the phenol consumption rate was not affected by the dissolved oxygen concentration. Thus, in order to reduce the model complexity it was assumed that, within the tested conditions, both productive and unproductive reaction rates were not affected by the dissolved oxygen concentration. Under these assumptions, the productive path was represented as follows:



To represent the unproductive path, the following reactions were proposed:



To express the phenol consumption rate (R_s) as a function of S , a steady-state for E_{ox} , $E_{ox}S$, $E_{red}S$ and E_{red} was assumed. Under this assumption the phenol consumption rate (R_s) is:

$$-\frac{dS}{dt} = k_p (E_{red}S) = R_s \quad (10)$$

Considering Eqs. (3) and (4), the oxygen consumption rate (R_{O_2}) is

$$R_{O_2} = R_s + \frac{1}{2}R_{unp} = k_p (E_{red}S) + \frac{1}{2}k_{unp} (E_{red}) \quad (11)$$

To further simplify the model, it was assumed that NADPH and NADP⁺ concentrations are nearly constant during the phenol metabolism. Because NADPH and NADP⁺ are intracellular metabolites, their total concentration ($N_T = N_{red} + N_{ox}$) is proportional to the biomass concentration (X). If the initial phenol concentration (S_0) is much lower than N_T , then both NADPH and NADP⁺ concentrations can be considered as constants. Additionally, if f_{red} is defined as the ratio N_{red}/N_T , then f_{red} is a constant also. Under this assumption, the following expression can be obtained:

$$R_s = a \left(\frac{bS + S^2}{c + dS + S^2} \right) X \quad (12)$$

where the expressions corresponding to the constants a , b , d and d are the following:

$$a = k_p q_E \left\{ \frac{N_T k_3 f_{red}}{k_p + N_T [k_4 (1 - f_{red}) + k_3 f_{red}]} \right\} \quad (13)$$

$$b = \frac{k_{unp}}{k_6} \quad (14)$$

$$c = \frac{k_{unp}}{k_1 k_6} \left\{ \frac{k_2 (k_p + k_5) + N_T [k_2 k_4 (1 - f_{red}) + k_3 f_{red} (k_p + k_5)]}{k_p + N_T [k_4 (1 - f_{red}) + k_3 f_{red}]} \right\} \quad (15)$$

According to the proposed model, the rate of the unproductive path is:

$$R_{unp} = \left(\frac{be}{S+b} \right) R_s \quad (17)$$

where:

$$e = \frac{k_s}{k_p} \quad (18)$$

The overall oxygen uptake rate during the phenol oxidation (R_{O_2}) due to productive and unproductive paths can be calculated by combining Eqs. (11), (12) and (17):

$$R_{O_2} = \left[\frac{S+b(1+e/2)}{S+b} \right] R_s \quad (19)$$

Using software package Gepasi 3, Eqs. (12) and (19) were fitted to the measured R_{O_2} values as a function of time, corresponding to the different initial phenol concentrations; obtained results are shown in Table 1. Figures 4 and 5 demonstrate that the proposed model adequately represents R_{O_2} and R_s as a function of time for all of the tested initial phenol concentrations. Using the fitted model, R_{O_2} profiles were calculated for different S_0 values; then, maximum oxygen consumed (OC_m) was calculated using Eq. (2) and the overall oxidation coefficient ($Y_{O/S}$) calculated as the ratio OC_m/S_0 . Figure 5b shows that $Y_{O/S}$ values obtained from the model were close to the experimental ones, confirming the validity of the developed model.

The fitted model was used to represent the total phenol concentration (TPh) and oxygen uptake rate (R_{O_2}) profiles during the fill and draw cycle shown in Fig. 1. In this case, the proposed model adequately predicts the linear decrease of TPh up to its depletion (Fig. 1a). Moreover, Fig. 1b shows that R_{O_2} increased as a function of time up to the substrate depletion at $t = 2.2$ h. Taking into account that these data were not used to fit the model, the agreement shown in Fig. 1 provides a validation of the proposed model.

Analysis of the developed model and comparison with other authors

Several conclusions arise from the analysis of the proposed model. The first one is that the kinetic coefficients obtained from respirometric data (e.g., R_{O_2} as a function of S) characterise the substrate consumption kinetic only if the oxidation coefficient is a constant. Note that the term between brackets in Eq. (19) corresponds to the instantaneous oxidation coefficient, which in fact varies as a function of the actual substrate concentration. For this reason, the overall oxidation coefficient was also a function of the initial phenol concentration.

According to the proposed model, dependences between R_{O_2} and R_s with S are quite different. While R_{O_2} as a function of S is close to the Haldane model, a Monod-type equation is more suited to describe the dependency of R_s with S (Fig. 5a). In most of the proposed models in the literature, the Haldane equation is an assumption to describe, with more or less accuracy, the biomass growth, phenol and oxygen consumption rates as a function of substrate concentration (Ben-Youssef and Vazquez-Rodriguez, 2011; Christen et al., 2012; Pramparo et al., 2012). Conversely, in the model proposed in the present work the inhibition of the respiration rate for high phenol concentrations arises as a natural consequence of the unproductive path during phenol metabolism (Eqs. (8), (9)). Indeed, in the case of an extremely stable $E_{red}S$ complex (e.g., $k_5 \cong 0$ or $k_6 \rightarrow \infty$), or if the free reduced enzyme (E_{red}) could not be oxidised by oxygen (e.g., $k_{unp} \cong 0$), coefficients

Coefficient	Units	Value	Standard deviation
a	Mmol $TPh \cdot g^{-1} TSS \cdot h^{-1}$	0.687	0.003
b	mM	2.00	0.03
c	mM ²	0.0205	0.0004
d	mM	1.91	0.02
e	--	2.92	0.02

b and c tend to zero (Eqs. (14), (15)), and in this case Eq. (19) of the proposed model can be simplified to obtain the Monod equation.

Several authors have found that the Haldane equation is not always suitable to represent the dependence of growth and substrate consumption rates as a function of S . For example, Marrot et al. (2006) studied the growth of activated sludge in phenol as a sole carbon source; these authors found that the specific growth rate increases with phenol concentration and then declines as substrate inhibition became important, in agreement with the Haldane equation. However, according to the data reported by the authors of the abovementioned work, the specific growth rate remains almost constant, about 10% of the observed maximum, for phenol concentrations higher than 10 mM. This behaviour is in agreement with the proposed model; Eqs. (12) and (19) show that both R_s and R_{O_2} tend to constant values at high phenol concentrations.

Feitkenhauer et al. (2001) proposed the following equation for the growth rate (R_x) of *Bacillus themoleovorans* as a function of phenol concentration:

$$R_x = \mu_{max} \left(\frac{S}{K_s + S + S^2/K_i} \right) (1 + S/K) X \quad (20)$$

This equation is similar to the proposed model; rearranging Eq. (16) the following expression can be obtained:

$$R_s = \left(\frac{ab}{d} \right) \left(\frac{S}{c/d + S + S^2/d} \right) (1 + S/b) X \quad (21)$$

From the comparison between Eqs. (20) and (21), it can be concluded that $K = b$, $K_s = c/d$ and $K_i = d$; thus, Feitkenhauer et al. (2001) found an empirical equation that is similar to the model developed in the present work.

Several authors have also reported that, for low phenol concentrations, the Monod model with a constant oxidation coefficient ($Y_{O/S}$) is enough to represent oxygen and phenol consumption kinetics (Chudoba et al., 1985; Orupold et al., 2001; Contreras et al., 2008; Lepik and Tenno, 2011). Under this condition, quadratic terms in Eq. (12) can be neglected, obtaining the following expressions:

$$R_s = \frac{ab}{d} \left(\frac{S}{c/d + S} \right) X \quad (22)$$

$$R_{O_2} = (1 + e/2) R_s \quad (23)$$

Equations (22) and (23) demonstrate that the proposed model tends to the Monod model with a constant oxidation coefficient ($Y_{O/S}$) at low phenol concentrations. Figure 5b shows that $Y_{O/S}$ values calculated by using the proposed model at low phenol concentrations are close to the reported values of other authors. Besides, according to Eq. (22), and using the constants shown in Table 1, the phenol saturation constant (K_s) can be calculated as the ratio $c/d = 10 \mu M$. This value is in accordance with

K_s values reported by other authors who have studied phenol biodegradation using respirometric techniques. Chudoba et al. (1985) reported a K_s of 14 μM ; Orupold et al. (2001) found K_s values ranging from 8 to 16 μM , corresponding to an activated sludge acclimated to phenolic compounds. Contreras et al. (2008) reported a value for $K_s = 11 \mu\text{M}$, while Lepik and Tenno (2011) informed $K_s = 4 \mu\text{M}$. These results demonstrate that the proposed model not only adequately represents the experimental results of the present paper, but also several findings reported by other authors.

CONCLUSIONS

This work demonstrates that while the phenol removal rate by acclimated activated sludge follows a Monod type kinetics, the oxygen uptake rate obeys a Haldane-type dependence with respect to the phenol concentration; thus, the phenol oxidation coefficient cannot be considered as a constant. A mathematical model based on a simplified version of the catalytic mechanism of the enzyme PH2MO was developed. The proposed model not only adequately represents the experimental results of the present paper, but also predicts several features regarding the aerobic phenol biodegradation that has been reported by other authors. The proposed model can be useful for predicting transient responses, such as substrate and DO concentrations as a function of time, in bioreactors experiencing an overload of phenolic wastewaters.

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