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Résumé de l'article

La dégradation photocatalytique de deux herbicides, le monolinuron (MLN) et le linuron (LN), a été étudiée dans une suspension de TiO₂ au contact de lumière solaire simulée. L'objectif de cette étude est de comparer leur photoréactivité et d'évaluer l'influence de différents paramètres tels que la concentration initiale en pesticide, la concentration en catalyseur et le flux photonique sur la cinétique de dégradation du monolinuron. L'étude comparative de la dégradation des deux herbicides a montré que les deux composés présentent une réactivité comparable avec le système TiO₂/lumière simulée. Dans les conditions opératoires de cette étude, la dégradation photocatalytique du LN et du MLN suit une cinétique de pseudo ordre 1. La pseudo-constante k_{obs} est inversement proportionnelle à la concentration initiale en pesticide et peut être décrite par une équation de Langmuir-Hinshelwood. L'étude de l'influence de la concentration en TiO₂ a permis de montrer que la cinétique de dégradation du MLN augmente avec la dose de catalyseur, mais un surdosage n'est pas nécessaire pour augmenter l'efficacité du procédé. Cette étude a permis également de montrer que la pseudo-constante d'ordre 1 de dégradation du MLN augmente linéairement avec le flux lumineux jusqu'à 580 W m⁻² puis décroît. Dans ces conditions, la recombinaison des électrons trous est favorisée. Enfin, la cinétique de dégradation du MLN en fonction du pH passe par un optimum de pH proche du pH de point de charge nulle (pH = 6).

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PHOTOCATALYTIC DEGRADATION OF MONOLINURON AND LINURON IN AN AQUEOUS SUSPENSION OF TITANIUM DIOXIDE UNDER SIMULATED SOLAR IRRADIATION

Dégradation photocatalytique du monolinuron et du linuron dans une suspension aqueuse de dioxyde de titane au contact de lumière solaire simulée

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ABSTRACT

The photocatalytic degradation of two phenylurea herbicides, monolinuron (MLN) and linuron (LN), was investigated in an aqueous suspension of TiO₂ using simulated solar irradiation. The objective of the study was to compare their photocatalytic reactivity and to assess the influence of various parameters such as initial pesticide concentration, catalyst concentration and photonic flux on the photocatalytic degradation rate of MLN and LN. A comparative study of the photocatalytic degradation kinetics of both herbicides showed that these two compounds have a comparable reactivity with TiO₂/simulated sun light. Under the operating conditions of this study, the photocatalytic degradation of MLN and LN followed pseudo first-order decay kinetics. The k_{obs} values indicated an inverse dependence on the initial herbicide concentration and were fitted to the Langmuir-Hinshelwood equation. Photocatalytic degradation rates increased with TiO₂ dosage, but overdoses did not necessarily increase the photocatalytic efficiency. The degradation rate of MLN increased with radiant flux until an optimum at 580 W m⁻² was reached and then decreased. Under these conditions,

an electron-hole recombination was favored. Finally, the photocatalytic degradation rate depended on pH, where an optimum was found at a pH value close to the pH of the point of zero charge (pH = 6).

Keywords: *Linuron; Monolinuron; phenyl urea herbicides; photocatalysis; titanium dioxide.*

RÉSUMÉ

La dégradation photocatalytique de deux herbicides, le monolinuron (MLN) et le linuron (LN), a été étudiée dans une suspension de TiO₂ au contact de lumière solaire simulée. L'objectif de cette étude est de comparer leur photoréactivité et d'évaluer l'influence de différents paramètres tels que la concentration initiale en pesticide, la concentration en catalyseur et le flux photonique sur la cinétique de dégradation du monolinuron. L'étude comparative de la dégradation des deux herbicides a montré que les deux composés présentent une réactivité comparable avec le système TiO₂/lumière simulée.

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Dans les conditions opératoires de cette étude, la dégradation photocatalytique du MLN et du LN suit une cinétique de pseudo ordre 1. La pseudo-constante k_{obs} est inversement proportionnelle à la concentration initiale en pesticide et peut être décrite par une équation de Langmuir-Hinshelwood. L'étude de l'influence de la concentration en TiO_2 a permis de montrer que la cinétique de dégradation du MLN augmente avec la dose de catalyseur, mais un surdosage n'est pas nécessaire pour augmenter l'efficacité du procédé. Cette étude a permis également de montrer que la pseudo-constante d'ordre 1 de dégradation du MLN augmente linéairement avec le flux lumineux jusqu'à 580 W m^{-2} puis décroît. Dans ces conditions, la recombinaison des électrons trous est favorisée. Enfin, la cinétique de dégradation du MLN en fonction du pH passe par un optimum de pH proche du pH de point de charge nulle (pH = 6).

Mots clés : Linuron; Monolinuron; herbicides phényl urées; photocatalyse; oxyde de titane

1. INTRODUCTION

In the last few decades, considerable attention has been given to the toxic and persistent pollutants coming from industrial and agricultural human activities. The application of photochemistry to environmental remediation and treatment has also been an active area of research and heterogeneous photocatalysis using UV/ TiO_2 system has often been proposed to eliminate these pollutants (BAHNEMANN *et al.*, 1994; CUNNINGHAM and SRIJARANAI, 1994; HERRMANN, 1999; MARINAS *et al.*, 2001; OLLIS and EL-EKABI, 1993). Titanium dioxide is widely used as photocatalyst because it is photochemically stable, non-toxic and low cost (AUGUGLIARO *et al.*, 1995; TAPOLOV *et al.*, 2000). This process is based on the generation of reactive species (electron/positive hole pairs) upon absorption of UV light ($\lambda < 368 \text{ nm}$) by the photocatalyst. In the presence of oxygen, the electron may induce reduction and leads to the formation of superoxide anions. Positive holes oxidize adsorbed organic substrates or react with water leading to the formation of hydroxyl radicals. These radicals are well known to be strong oxidant agents (the reduction potential is as high as 2.8 V/HSE) and react efficiently without any selectivity with most organic substrates (D'OLIVEIRA *et al.*, 1990; OKAMOTO *et al.*, 1985; OLLIS *et al.*, 1989).

Phenylurea herbicides were introduced in the 1950's and their use has considerably increased. Monolinuron (MLN) and Linuron (LN) (Figure 1) are widely used in pre- and post-emergence for treatment of both crop and non-crop cultures. These herbicides are moderately persistent in soils, with a field half-life of approximately 30 to 150 days and are slightly to

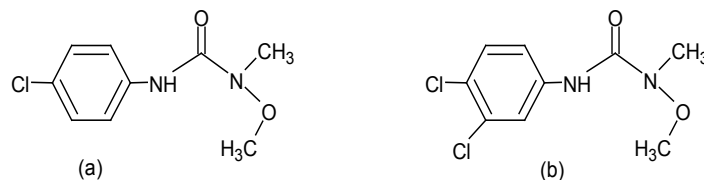


Figure 1. Chemical structure of (a) monolinuron (MLN) and (b) linuron (LN).

Formule chimique du (a) monolinuron (MLN) et (b) linuron (LN).

moderately soluble in water ($s = 735$ and 81 mg L^{-1} at 25°C for MLN and LN respectively) (TOMLIN, 2000). Consequently, MLN and LN can give rise to important residues in soil and in water.

The aim of the present work was to study the photocatalytic degradation of LN and MLN in presence of TiO_2 in aqueous solution using simulated solar irradiation. The effect of various parameters, such as herbicide concentration, photonic flux, mass of catalyst and pH on the decay of MLN, was investigated. These herbicides were the subject of direct photolysis but there are no published papers concerning their photocatalytic transformation.

2. MATERIALS AND METHODS

2.1 Reactants

All reagents were of the purest grade commercially available and were used without further purification. Monolinuron (3-(4-chlorophenyl)-1-methoxy-1-methylurea, 99%) was Riedel- de Haën and linuron (3-(3,4-dichlorophenyl)-1-methoxy-1-methylurea, 98%) was from Chem. Service. Titanium dioxide Degussa P25 was mostly in the anatase form (70% anatase, 30% rutile) with an average particle size of 30 nm and BET surface area of $50 \text{ m}^2 \text{ g}^{-1}$. Acetonitrile, HPLC grade product, was purchased from Merck. Current reagents such as NaHCO_3 and Na_2CO_3 were analytic grades (Merck). Water was purified by Milli-Q system (Millipore) and controlled by its resistivity ($18 \text{ M}\Omega \text{ cm}^{-1}$).

2.2 Photocatalytic experiments

Suspensions of TiO_2 ($0-1.5 \text{ g L}^{-1}$) in aqueous solutions of MLN or LN ($0.8-8 \text{ mg L}^{-1}$) were irradiated in a CPS+ ATLAS Suntest simulator, equipped with a Xenon lamp (irradiance between 250 and 765 Wm^{-2}) with a cut-off at $\lambda = 290 \text{ nm}$. The reactor was an open water-jacketed cylindrical vessel (Figure 2). Experiments were carried out under aerated



Figure 2. CPS+ ATLAS Suntest simulator.
Simulateur solaire CPS+ ATLAS Suntest.

conditions ($[O_2] = 7.4 \text{ mg L}^{-1}$) at a temperature of the irradiated solutions maintained at $17 \pm 3^\circ\text{C}$. In all experiments, the total irradiated solution volume was 250 mL with a path length of 4 cm. TiO_2 was maintained in suspension before and during irradiation by magnetic stirring. Aliquots of suspensions were sampled at the end of the adsorption for a period of one hour and during irradiation at selected time intervals. Samples were filtered through hydrophilic polypropylene filters ($0.45 \mu\text{m}$) to remove TiO_2 particles prior to HPLC analysis.

2.3 Analysis

UV-visible spectra were recorded on Unicam HELIOS α double beam spectrophotometer. The disappearance of substrates and formation of products were monitored by HPLC on a WATERS chromatograph equipped with two pumps (WATERS 515) and a photodiode array detector (WATERS 996). The column was an Interchim UP3* 12QK ($3 \mu\text{m}$, $125 \text{ mm} \times 4 \text{ mm}$). The eluant was a mixture of 40/60 (V/V) acetonitrile-water + 1‰ acetic acid for MLN and 50/50 (V/V) acetonitrile-water for LN, with a flow rate of 0.7 mL min^{-1} and detection wavelength at 244 nm. Chloride ion concentrations were determined by ionic chromatography. A cartridge glass IC column (Metrosep Anion Dual 1: $150 \times 3.0 \text{ mm}$) was used. The eluant was a mixture of Na_2CO_3 (2.5 mmol L^{-1}) and NaHCO_3 (2.4 mmol L^{-1}). Under these conditions, retention times of LN, MLN and chloride ions were 10.3, 10.5 and 5.5 minutes respectively.

3. RESULTS AND DISCUSSION

3.1 UV spectra

The UV absorption spectra of monolinuron and linuron in aqueous solution are given in Figure 3. These two herbicides have similar absorption spectra with a maximum absorption band located at 243 nm and 246 nm, and with a molar absorption coefficient of $17,500 \pm 50 \text{ L mol}^{-1} \text{ cm}^{-1}$ and $16,500 \pm 50 \text{ L mol}^{-1} \text{ cm}^{-1}$ for MLN and LN respectively. The

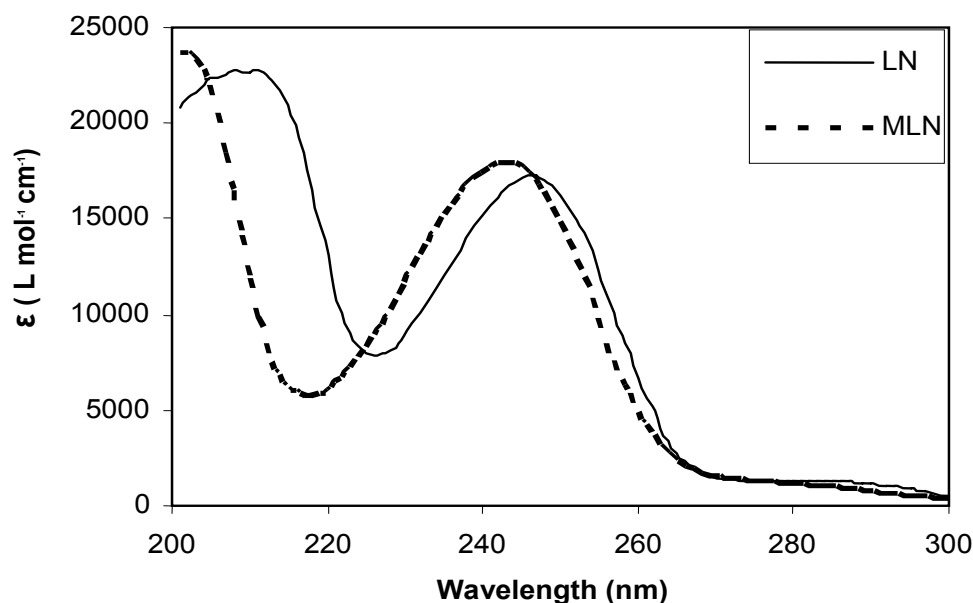


Figure 3. UV spectra of linuron and monolinuron.
Spectres UV du linuron et du monolinuron.

weak bathochromic shift ($\Delta\lambda = 3$ nm) and hypochrom effect ($\Delta\varepsilon = 1000$ L mol⁻¹ cm⁻¹) are due to the presence of a second chlorine atom on the benzene ring for LN. For both pesticides, a shoulder with a maximum around 280 nm is also observed with a molar absorption coefficient of 1,200 L mol⁻¹ cm⁻¹.

3.2 Direct photolysis and photocatalysis

In the absence of TiO₂, no significant decrease in the concentration of both herbicides was observed under simulated solar irradiation for about two hours (Figure 4). In this case, the amount of pesticide removed is less than 8% for an incident photonic flux of 250 W m⁻². The very low molar absorption coefficient for both molecules above 300 nm associated with their weak quantum yield, result in a good photostability of these pesticides under sunlight irradiation, which implies that these compounds would be removed very slowly by direct photolysis. In presence of TiO₂, amounts of MLN and LN adsorbed over photocatalyst in the dark were less than 5%, thus degradation of these herbicides is exclusively due to photocatalytic processes via the formation of hydroxyl radicals generated at the surface of the semiconductor (NARGIELLO and HERZ, 1993; OLLIS *et al.*, 1989).

For our experimental conditions ([Herbicide] = 2 · 10⁻⁵ mol L⁻¹; [TiO₂] = 0.2 g L⁻¹; photonic flux P = 250 W m⁻²), data are in a good agreement with a pseudo-first-order reaction, as evidenced by plotting ln(C_t/C₀) versus irradiation time (insert in Figure 4). The apparent rate constants calculated for monolinuron and linuron irradiated in separate solutions are 0.081 min⁻¹ and 0.082 min⁻¹ with a half-life time of about 8.5 min indicating that MLN and LN have a similar reactivity with TiO₂/Simulated sunlight. The substitution of the benzene ring with one or two Cl atoms does not affect the photocatalytic reactivity of these pesticides.

In order to determine whether MLN and LN were completely mineralized, the concentration of chloride ions and TOC (Total Organic Carbon) was monitored at different irradiation times.

As shown in Figures 5a and 5b, total dechlorination of both pesticides is obtained after the corresponding time of illumination required for the total disappearance of the initial substrate. On the other hand, TOC removal is only 25% for LN and MLN. This means that mineralization of both substrates

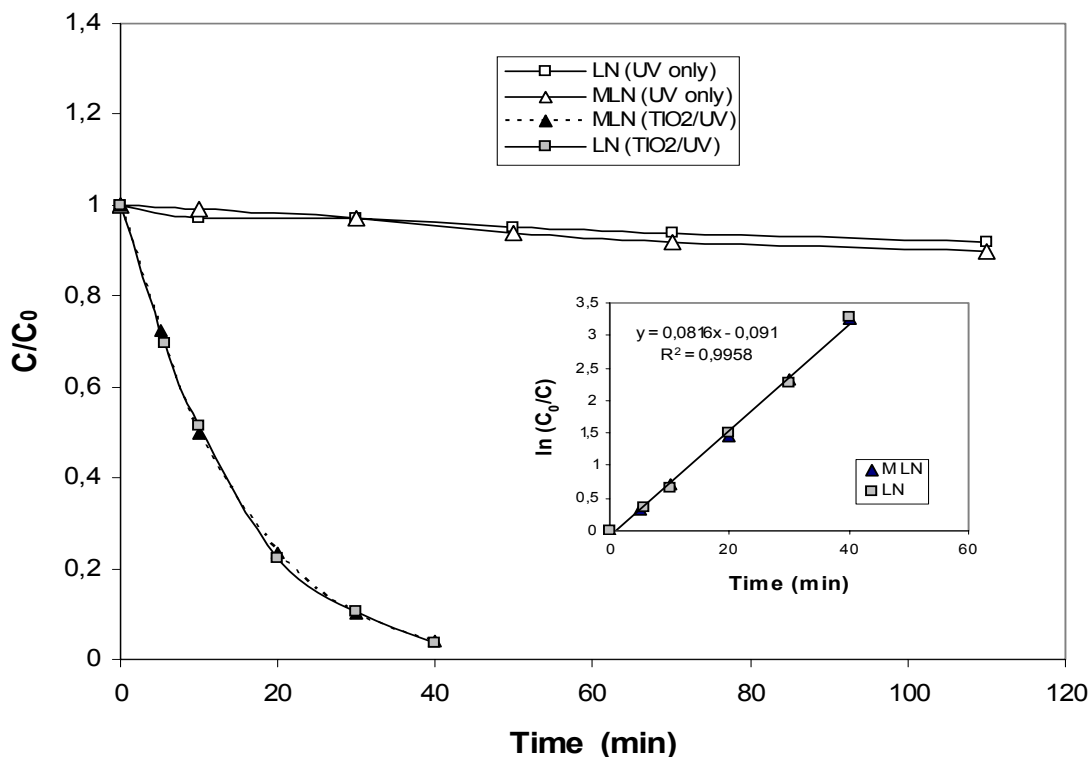


Figure 4. Kinetics of the disappearance of MLN and LN ($C_0 = 2 \cdot 10^{-5}$ mol L⁻¹) in the presence of 0.2 g L⁻¹ of TiO₂ at 250 W m⁻² (insert: $\ln(C_t/C_0) = f(\text{time})$).
Cinétique de disparition du MLN et LN ($C_0 = 2 \cdot 10^{-5}$ mol L⁻¹) en présence de 0,2 g L⁻¹ de TiO₂ à 250 W m⁻² (insertion : $\ln(C_t/C_0) = f(\text{temps})$).

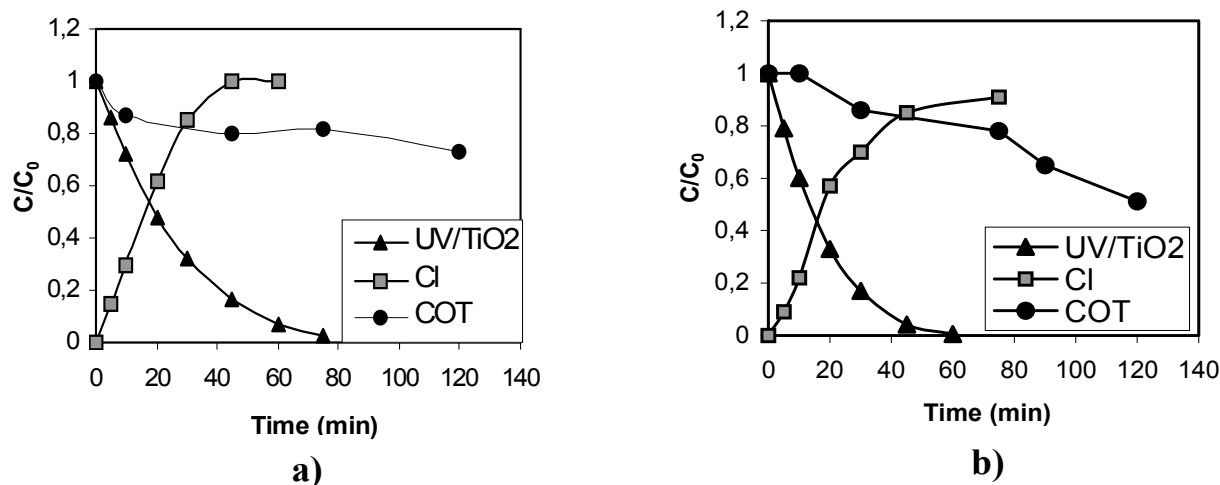
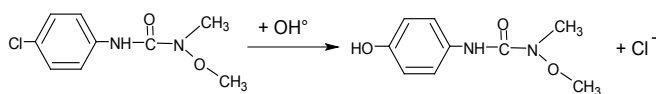


Figure 5. Kinetics of the disappearance of the two herbicides, appearance of chloride ions and evolution of the TOC ($C_0 = 2 \cdot 10^{-5} \text{ mol L}^{-1}$, $[\text{TiO}_2] = 0.2 \text{ g L}^{-1}$, $P = 250 \text{ W m}^{-2}$) - (a) linuron, (b) monolinuron.
Cinétique de disparition des deux herbicides, apparition des ions chlorure et évolution du COT ($C_0 = 2 \cdot 10^{-5} \text{ mol L}^{-1}$, $[\text{TiO}_2] = 0,2 \text{ g L}^{-1}$, $P = 250 \text{ W m}^{-2}$) - (a) linuron, (b) monolinuron.

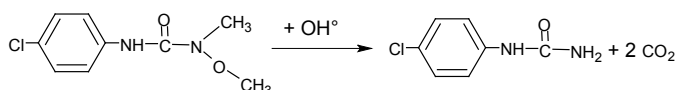
is slow in spite of their photodegradation. Chromatographic analysis of MLN solution ($C_0 = 3.7 \cdot 10^{-5} \text{ mol L}^{-1}$), irradiated during 45 min, in the presence of 0.2 g L^{-1} of TiO_2 , shows the formation of two major photoproducts (P_1 and P_2) (Figure 6). UV-visible spectra analysis of these products indicates that they absorb mainly at the same wavelength range as MLN, confirming that primary OH radical attack does not destroy the aromatic ring. The kinetics of formation and disappearance of P_1 and P_2 show that they are primary products.

Identification of these compounds was not investigated, but according to (i) our kinetic results for chloride ion formation and TOC removal and (ii) the degradation scheme of similar phenyl urea herbicides such as metobromuron (AMINE-KHODJA *et al.*, 2005) or monuron (PRAMAURO and VINCENTI, 1993) reported in the literature, P_1 and P_2 could result from the simultaneous attack of radical species on two positions of the initial molecule with:

Hydroxylation of the aromatic ring by substitution of Cl atom.



An attack of the alkyl urea group leading to the partial mineralization of initial substrate.



The kinetic behaviour of these derivatives (Figure 6) clearly confirms that these aromatic compounds undergo further transformation, probably into non-aromatic products which cannot be detected by HPLC analysis under the experimental conditions used in this study. Total mineralization of these intermediates will require a long irradiation time since residual TOC still remains after two hours of irradiation.

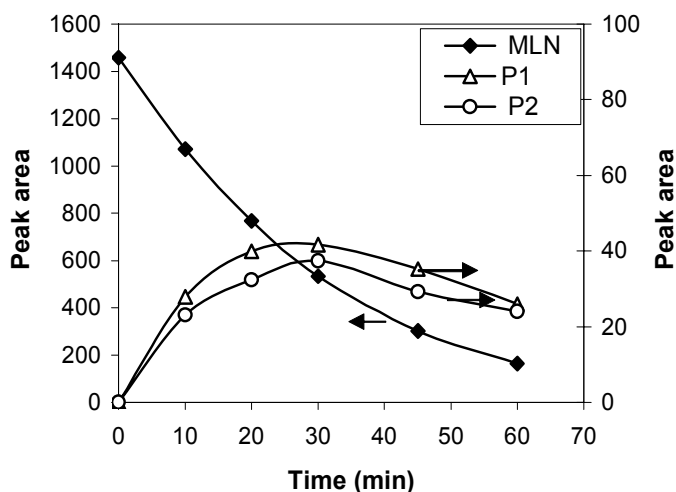


Figure 6. Kinetics of the disappearance of MLN and formation of intermediates ($[\text{MLN}] = 3.7 \cdot 10^{-5} \text{ mol L}^{-1}$, $[\text{TiO}_2] = 0.2 \text{ g L}^{-1}$, $P = 250 \text{ W m}^{-2}$).
Cinétique de disparition du MLN et formation des intermédiaires ($[\text{MLN}] = 3,7 \cdot 10^{-5} \text{ mol L}^{-1}$, $[\text{TiO}_2] = 0,2 \text{ g L}^{-1}$, $P = 250 \text{ W m}^{-2}$).

3.3 Effect of various parameters on monolinuron photocatalytic degradation

Initial herbicide, TiO_2 concentrations and photonic flux are important parameters which can affect photocatalytic degradation rates. The effect of these three parameters has been studied at natural pH (pH = 5), on the photocatalytic degradation of monolinuron (MLN) chosen as the pesticide. For all experiment sets, the data found for the degradation of MLN are in good agreement with a pseudo-first-order reaction with respect to concentrations in the experimental range ($0.98 < r^2 < 0.99$).

$$r_{\text{MLN}} = -\frac{d[\text{MLN}]}{dt} = k_{\text{obs}}[\text{MLN}] \quad (1)$$

3.3.1 Effect of TiO_2 concentration

Figure 7 shows the influence of TiO_2 concentration ($[\text{TiO}_2] = 0.1$ to 1.5 g L^{-1}) on the degradation rate of MLN at 4 mg L^{-1} and for a photonic flux of 250 W m^{-2} . As expected, the apparent rate constant (k_{obs}) was found to increase with the increasing concentration of semiconductor. This enhancement is due to the increase of the number of active sites with the TiO_2 concentration. However, k_{obs} tends to a limit that corresponds

to the total absorption of the photonic flux by the catalyst as illustrated in Figure 7 (average absorbance between 290 and 400 nm of the TiO_2 suspension for 4 cm path length versus k_{obs}). For our experimental conditions, this limit seems to be reached at 1 g L^{-1} for which 99% of the light is absorbed. Above this concentration, a plateau is observed as it is mentioned by several authors (INEL and OKTE, 1996; Le CAMPION *et al.*, 1999; PARRA *et al.*, 2002; WONG and CHU, 2003). Figure 7 also points out that at low TiO_2 concentration ($< 0.1 \text{ g L}^{-1}$) or weak absorption, the degradation of monolinuron occurs at very low rate.

3.3.2 Effect of herbicide concentration

The effect of the monolinuron concentration has been studied in a range between 0.8 to 8 mg L^{-1} for a constant TiO_2 dosage ($[\text{TiO}_2] = 0.2 \text{ g L}^{-1}$) and a constant photonic flux ($P = 250 \text{ W m}^{-2}$). Figure 8 shows that the apparent first-order rate constant k_{obs} decreases with the increasing of the initial concentration of MLN. This can be explained by the competition of photoproducts with substrate for the adsorption on the active sites of the TiO_2 surface (OLLIS *et al.*, 1989; PRAMAURO and VINCENZI, 1993).

In the last few decades, the Langmuir-Hinshelwood expression has been successfully used to describe the dependence of the reaction rate r_{MLN} on the initial concentration of the

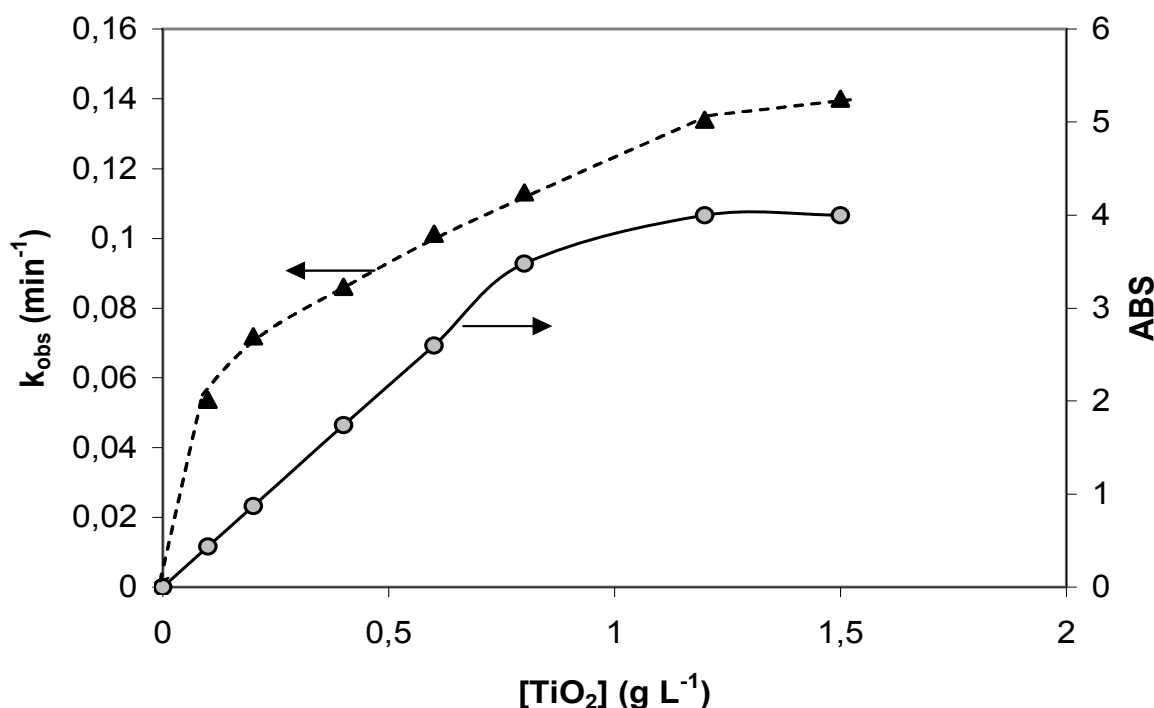


Figure 7. Effect of TiO_2 concentration on the pseudo-first-order rate constant.
Influence de la concentration en TiO_2 sur la pseudo-constante d'ordre 1.

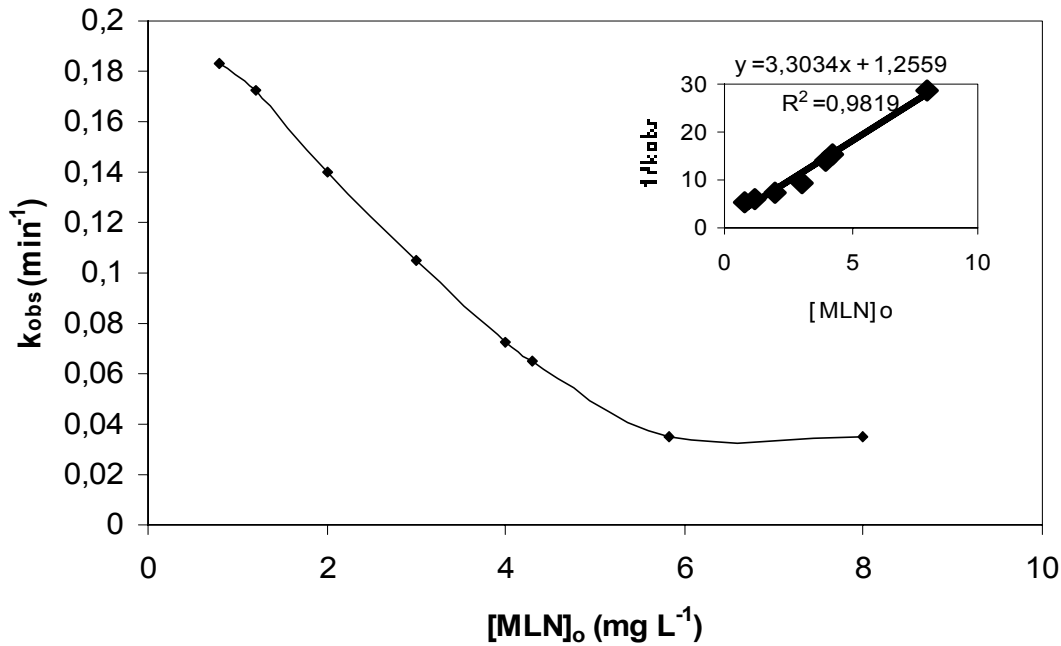


Figure 8. Effect of initial MLN concentration on the pseudo-first-order rate constant.
Influence de la concentration initiale en MLN sur la pseudo-constante d'ordre 1.

organic substrate $[\text{MLN}]_o$ for heterogeneous photocatalytic degradation (HEREDIA *et al.*, 2001):

$$r_{\text{MLN}} = -\frac{d[\text{MLN}]}{dt} = k_{\text{obs}}[\text{MLN}] = \frac{kK_{\text{MLN}}[\text{MLN}]}{1 + K_{\text{MLN}}[\text{MLN}]_o} \quad (2)$$

where: K_{MLN} is the Langmuir-Hinshelwood adsorption equilibrium constant of monolinuron and k is the rate constant at the surface reaction. A plot of $1/k_{\text{obs}}$ vs. initial monolinuron concentration $[\text{MLN}]_o$ gave a straight line (Figure 8) indicating that the kinetics fit with a Langmuir-Hinshelwood equation :

$$\frac{1}{k_{\text{obs}}} = \frac{1}{kK_{\text{MLN}}} + \frac{[\text{MLN}]_o}{k} \quad (3)$$

From the intercept and the slope, k and K_{MLN} can be easily calculated. The results obtained are $k = 0.30 \text{ mg L}^{-1} \text{ min}^{-1}$ ($1.4 \cdot 10^{-6} \text{ mol L}^{-1} \text{ min}^{-1}$) and $K_{\text{MLN}} = 2.6 \text{ L mg}^{-1}$ ($5.6 \cdot 10^5 \text{ L mol}^{-1}$). The determination of these constants for the photocatalytic transformation of monolinuron has no absolute meaning because they depend on various parameters such as TiO_2 concentration, pH, photonic flux, etc. However, the good correlation obtained from the plot $1/k_{\text{obs}}$ vs. initial monolinuron concentration $[\text{MLN}]_o$ indicates that the photocatalytic degradation of MLN occurred mainly at the

surface of the semiconductor. As a consequence, the kinetics depend on the equilibrium of sorption / desorption of the pesticide, and thus on its initial concentration.

3.3.3 Effect of radiant flux

The incident flux, which is of great importance in the photocatalytic process, has been the subject of many studies. OLLIS *et al.* (1989), D'OLIVEIRA *et al.* (1990) and ALSAYYED *et al.* (1991) reported that at low intensity levels of illumination, the degradation rate is of first-order in intensity. However, at high intensity, the reaction rate increases with the square root of the intensity level. INEL and OKTE (1996) also found a similar rate shift from first order to half order. The initial degradation rate of 4-nitrophenol versus radiant flux was also investigated in the range of 0-180 W m^{-2} (CHEN and RAY, 1998), where the effect was correlated by: $k \propto P^{0.84}$. This was explained by the fact that at low levels of intensity the degradation rate is neither catalyst dependent nor mass transfer dominated.

For our study, the effect of the radiant flux on the photocatalytic degradation of MLN is shown in Figure 9 for a range varying from 250 to 765 W m^{-2} [TiO_2] = 0.2 g L^{-1} , $[\text{MLN}]_o = 4 \text{ mg L}^{-1}$). It can be noted that the pseudo-rate constant passes through a maximum with an optimum at $P_{\text{max}} = 580 \text{ W m}^{-2}$. At low fluxes ($P < 580 \text{ W m}^{-2}$), the reaction rate is proportional to the radiant flux, indicating that the process works in a good photocatalytic regime: the incident photons are efficiently converted into actives species, corresponding to the degradation mechanism described above

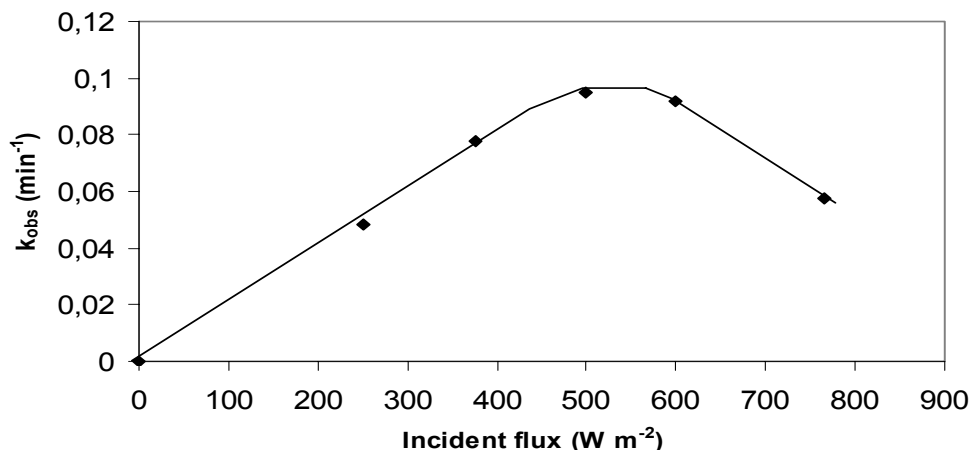
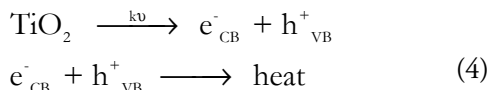


Figure 9. Effect of radiant flux on the pseudo-first-order rate constant.
Influence du flux lumineux sur la pseudo-constante d'ordre 1.

(HERRMANN, 1999; VULLIET *et al.*, 2003). At high fluxes, the rate decreases with the increase of radiant flux, indicating an increase of the recombination of photoproduced charges relative to the oxidation of substrate at high intensities, as has been reported elsewhere (Al-SAYYED *et al.*, 1991; D'OLIVEIRA *et al.*, 1990; HERRMANN, 1999; VINCZE and KEMP, 1990):



3.3.4 Effect of initial pH

An important parameter that affects the rate of MLN degradation is the pH of the suspension. When substrates are dissolved in water, the effect consists in the change in:

- adsorption characteristics of molecules depending on their polarity and the charge of the TiO_2 surface at different pH levels;
- predominance of radical species depending upon the hydroxyl or proton ion concentration.

It is well known that the point of zero charge (pzc) for TiO_2 is at a pH between 5.6 and 6.4 and the surface coated with hydroxyl groups may add or release protons, as shown in the following equilibrium (CHEN and RAY, 1998; TERZIAN and SERPONE, 1995):



In this study, the pH was adjusted in acidic media with HCl or HClO_4 and in basic media with NaOH or NH_3 . The effect of the addition of these bases and acids on the photocatalytic degradation rate of MLN ($C_0 = 4 \text{ mg L}^{-1}$; $\text{TiO}_2 = 0.2 \text{ g L}^{-1}$; $P = 250 \text{ W m}^{-2}$) is illustrated in Figure 10. Results indicate that pH value has a significant effect at both low and high pH:

- At $\text{pH} < 6.3$, the photocatalytic degradation rate of MLN depends on the acid used. When HCl is used, the degradation rate decreases with pH. This decrease can be explained by (i) the low OH^- concentration which hinders the formation of OH^\bullet and subsequently reduces the degradation rate and (ii) an inhibition of photocatalytic reactions with chloride ions which adsorb on the surface of TiO_2 and compete with MLN on the adsorption sites. At a pH around 5, the inhibition of Cl^- is small compared to the degradation rate obtained in water (Figure 10), but this inhibition increases with Cl^- concentration particularly at low pH values. Similar results were reported for the degradation of nitrophenols by AUGUGLIARO *et al.* (1991) and by CHEN and RAY (1998). When HClO_4 is used, the degradation rate increases with perchlorate ion concentration which cancels the effect of pH observed with HCl. Similar results were obtained by IRMAK *et al.* (2004) who showed that oxyhalogens such as IO_4^- , BrO_3^- , ClO_3^- increase the degradation rate of organic pollutants by capturing the electrons ejected from TiO_2 so that the probability of recombination of e^-/h^+ pairs will decrease.

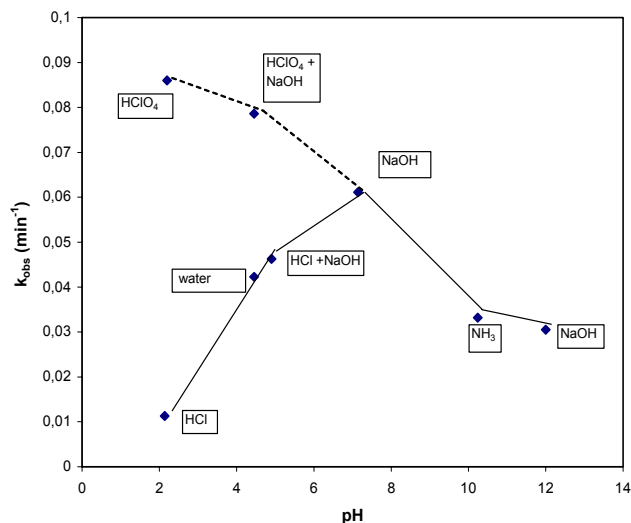


Figure 10. Effect of pH on the pseudo-first-order rate constant.

Influence du pH sur la pseudo-constante d'ordre 1.

- At $\text{pH} > 6.3$, the degradation rate decreased with the increasing pH value beyond the pzc. In this case, both bases used have a low influence on the rate constant. In this range of pH, the high OH^- ion content of the system enhances the electron-holes separation. However further increments of pH would retard the rates. This can be explained by the effect of charge repulsion between TiO_2 particles (negatively charged) and MLN molecules or hydroxide ions. Repulsion between TiO^- species present on the TiO_2 surface and MLN would occur through more atoms with negative partial charges localized on the nitrogen and oxygen of the urea function (Figure 11). These charges were calculated using hyperchem 6.0 software by the AM1 method.

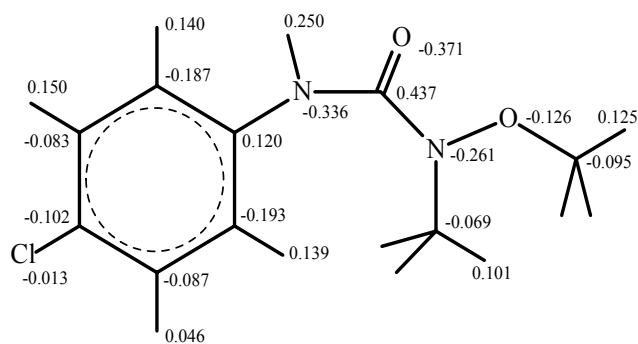


Figure 11. Charge densities on monolinuron atoms.
Densité de charge sur les atomes du monolinuron.

4. CONCLUSION

The photocatalytic degradation of two phenylurea herbicides (LN and MLN) in aqueous solution has been studied in the presence of TiO_2 Degussa P-25 under simulated solar irradiation. Results show that photocatalytic degradation of both herbicides follows pseudo-first-order decay. The kinetics of degradation of MLN and LN are very similar even though the benzene ring is substituted with 2 Cl atoms. This study pointed out the dependence of the photocatalytic rates on operating conditions such as TiO_2 dosage, herbicide concentration, photonic flux and pH, which are mainly related to the reactions at the interface of the catalyst such as adsorption / desorption equilibrium, electrostatic interaction and activation of the catalyst surface with light.

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