



PT-1-48

## KINETICS AND ADSORPTION COMPARATIVE STUDY ON PHOTOCATALYTIC DEGRADATION OF o-, m- AND p-CRESOL

*E. Pulido Melián\*, O. González Díaz, J. Araña and J. M. Doña Rodríguez*

Fotocatálisis y Electroquímica Aplicada al Medio-Ambiente (FEAM). Unidad Asociada al CSIC. CIDIA (Depto. de Química), Edificio del Parque Científico Tecnológico, Campus Universitario de Tafira, 35017, Las Palmas, España. UNIVERSIDAD DE LAS PALMAS DE GRAN CANARIA. Tlf. +34-928-45-72-99, Fax + 34-928-45-73.97. Email: [elisendapm80@hotmail.com](mailto:elisendapm80@hotmail.com)

Photocatalytic degradations with TiO<sub>2</sub> at different concentrations (10-125ppm) of o-, m-, and p- cresol have been studied. The initial degradation rates between described concentration range of these compounds are nearly the same for each substance (maxima differences being  $0.37 \cdot 10^{-7} \text{ M}\cdot\text{s}^{-1}$  for m-cresol,  $0.41 \cdot 10^{-7} \text{ M}\cdot\text{s}^{-1}$  for o-cresol,  $0.17 \cdot 10^{-7} \text{ M}\cdot\text{s}^{-1}$  for p-cresol). We can observe a clear maximum of initial degradation rate for m-cresol y o-cresol at 50 ppm. In contrast, p-cresol presents little variations along described concentration range without any clear maximum.

Adsorption and rate constants have been obtained by Langmuir-Hinselwood equation from initial concentration ( $C_o$ ) and velocity values ( $r_o$ ):

$$r_o = \frac{k \cdot K \cdot C_o}{1 + K \cdot C_o}$$

For this model, the rate of reaction is proportional to the fraction of surface covered by the substrate. At first, this model was developed in order to describe solid-gas reactions. Nevertheless this model has been recently applied to solid-liquid systems.

Degradation for diluted solutions of cresol follows pseudo-first-order kinetics. Initial rates are so obtained from next product,

$$r_o = k \cdot K \cdot C = k^o \cdot C_o$$

where  $k^o$  are first-order pseudo-constants for each concentration.

FTIR studies show cresol isomers interaction with catalyst surface. The region between  $1550\text{-}1300 \text{ cm}^{-1}$  was selected to emphasize  $\nu\text{C}=\text{C}$  and  $\delta\text{OH}$  vibration changes after interactions. O-cresol and m-cresol are adsorbed on catalyst surface by hydrogen bridges. p-cresol interaction is a mixture between cresolate formation and H-bond, as we have observed for phenol.

Even little proportion of cresol isomers in solution suffers adsorption on catalyst surface, interaction fits well Freundlich and Langmuir models for a given concentration range. p-cresol presents clear different behaviour from the other two studied isomers. As we have described, FTIR studies agree with obtained adsorption Langmuir constants.



## References:

1. Erdal Kusvuran, Ali Sammil, Osman Malik Atanur, Oktay Erbatur. Photocatalytic degradation kinetics of di- and tri- substituted phenolic compounds in aqueous solution by  $\text{TiO}_2/\text{UV}$ . *Applied CatalysisB: Environmental* 58(2005)211-216.
2. S. Bekkouche, M. Bouhelassa, N. Hadj Salah, F.Z. Meghlaoui. Study of adsorption of phenol on titanium oxide( $\text{TiO}_2$ ). *Desalination*, 166(2004)355-362.
3. José Pedro S. Valente, Pedro M. Padilla, Ariovaldo O. Florentino. Studies on the adsorption and kinetics of photodegradation of a model compound for heterogeneous photocatalysis onto  $\text{TiO}_2$ . *Chemosphere* 2006.
4. Didier Robert, Sandra Parra, César Pulgarin, André Krzton, Jean Victro Weber. Chemisorption of phenols and acids on  $\text{TiO}_2$  surface. *Applied Surface Science* 167(2000) 51-58.
5. Vinod K. Gupta, C.K. Jain, Imran Ali, M. Sharma, V.K. Saini. Removal of cadmium and nickel from wastewater using bagasse fly ash- a sugar industry waste. *Water Research* 37(2003) 4038-4044.