



## BIODEGRADATION OF DICOFOL IN A DOUBLE CHAMBER MICROBIAL FUEL CELL

Pedro Perez-Rodriguez<sup>a</sup>, Jose A. Rodríguez-de la Garza<sup>a\*</sup>, Leopoldo J. Ríos-González<sup>a</sup>, Yolanda Garza-García<sup>a</sup>, Silvia Y. Martínez-Amador<sup>b</sup>

<sup>a</sup>Departamento de Biotecnología, Facultad de Ciencias Químicas, Universidad Autónoma de Coahuila. Blvd. Venustiano Carranza s/n Col República. Saltillo, Coahuila, C.P. 25280, México.

<sup>b</sup>Departamento de Botánica, Universidad Autónoma Agraria Antonio Narro. Calzada Antonio Narro 1923, Buenavista, Saltillo, Coahuila.

*Key words: Microbial fuel cells, dicofol, electricity.*

**Introduction.** Environmental pollution has been a global problem and causes certain dangers to human life. Organochlorine pesticides, dicofol, are well known for their toxicity, widespread occurrence and their bioaccumulative abilities in the environment [1].

Microbial fuel cells (MFC) have recently been used for successful bioremediation of a number of chemicals. An MFC is a device that uses microbes to convert the chemical energy stored in organic or inorganic compounds into electricity, providing a low-cost and low-maintenance energy as well as a process that produces very little sludge. MFC have been used as a method for either reductive transformation contaminants such as nitrate and perchlorate or for producing value-added products [2].

The aim of the present study was to assess the feasibility of a two chamber MFC to enhance biodegradation of dicofol.

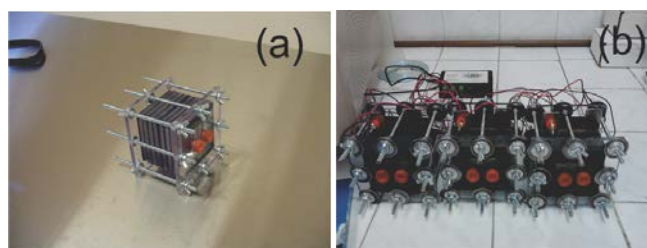
**Methods.** Electrodes (4 x 4 cm) were made out of stainless steel and coated with graphite. MFC was made of 6 mm polycarbonate plates (7 x 7 cm), neoprene gaskets were placed between each polycarbonate plate to avoid leakage from the anodic and cathodic chambers (fig 1). Voltage was recorded using a Madgetech Quadvolt datalogger connected to a computer. Mineral medium containing dicofol (40 ppm) and glucose as co-substrate was added to anodic chamber and sealed, the cathodic chamber was filled with  $K_3[Fe(CN)_6]$  (0.05 M). Anodic and cathodic chamber were separated by a protonic exchange membrane (Ultrax CMI 7000). Dicofol was quantified by GC ECD Varian Star 3400 CX equipped with a column CP-Sil CB for pesticides. Glucose consumption was determined by Chemical oxygen demand (COD) according to APHA standard methods.

**Results.** MFC with graphite coated electrode showed a better performance compared with the MFC with the stainless steel electrode. In table 1 its shown the biodegradation percentage for both cases, achieving 40.5 % for the case of the coated electrode MFC.

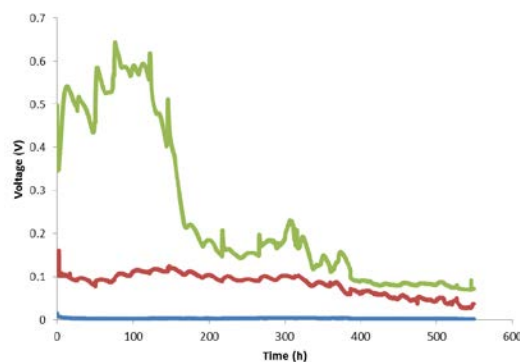
**Table 1.** Biodegradation of dicofol in a MFC using two types of electrodes in the anodic chamber.

MFC coated electrode	MFC stainless steel electrode
40.5 %	14.5 %

Voltage generation was higher in the case of the MFC with the graphite coated electrode with a maximum potential of 0.627 V. After 154 h potential decreased rapidly due to COD removal in the case of MFC with graphite coated electrode.



**Figure 1.** (a) MFC Assembly; (b) MFC used in the present study.



**Figure 2.** Voltage generation in a double chamber MFC for dicofol biodegradation. Green line MFC graphite coated electrode; red line stainless steel electrode, blue line control.

**Conclusions.** . The results obtained in the present study demonstrate the potential of MFC to enhance biodegradation of organic pollutant such as pesticides.

**Acknowledgements.** To de Mexican Council of Science and Technology (CONACYT) for providing scholarship to Perez-Rodriguez for carrying out this work as a part of his Master's thesis.

**References.**

1. Gregory, K.B., Bond, D.R., and Lovley, D.R., (2004). *Environ Microbiol.* 6 (1): 596–604.
2. Huang L., Chai X., Quana X., Logan B. E., Chenc G. (2012). *Biores Technol.* 111 (1): 167–174.