Composite Electrode using Graphite and Opuntia imbricata to Develop an Anodophilic Biofilm in a Microbial Fuel Cell


INTRODUCTION

Microbial Fuel Cells (MFC), are bioelectrochemical devices capable of converting energy stored in organic matter to electricity using bacteria as biocatalyst, and are rapidly gaining acceptance as an alternative energy technology (Ryckelynck et al., 2005; Picioreanu et al., 2008; Deng et al., 2010), due to its great versatility to use different substrates from renewable sources and convert them into harmless by-products with simultaneous production of electricity (Jiang et al., 2010). Production of electricity by MFC happens when bacteria extracts electrons from these substrates and migrate to the anode (negative terminal) in the anodic chamber and flow to the cathode (positive terminal) trough a conductive material and a resistor, or operated under a load (Reguera et al., 2006). These Electrons can be transferred to the anode by electron mediators or shuttles (Rabaey et al., 2005), by direct membrane associated electron transfer (Reguera et al., 2006), or conduction by nanowires (Rabaey et al., 2005) produced by the bacteria.

In the conduction mechanism, were bacteria transfer electrons to the anode from their membrane-bound electron carriers to the electrode via conductive materials in the biofilms matrix creating what has been called biofilms anode (Bond and Lovley, 2003; Picioreanu et al., 2010). The matrix of EPS of at least some anode-respiring biofilms contains highly conductive nanowires that extend for tens of micrometers (Reguera et al., 2006). In the model describe by it predicts how fast the anode biofilm grows, how fast substrate is oxidized and how much current is generated under different conditions. If the biofilm is too thin, there are insufficient bacteria to generate a substantial current, but if the biofilm is too thick, the current is reduced because electrons cannot reach the anode. In more conductive biofilms, other factors, such as substrate mass transport, become important (Jones, 2008; Liu et al., 2010). Cheng et al. (2008) suggested that controlling biofilm thickness, by using a turbulent environment in the microbial fuel cell to regulate the detachment of cells from the biofilm, might be one method of maximizing the current that is produced (Rodriguez-Martinez et al., 2005; Cheng et al., 2008).

In the present study, a two-chamber MFC was used, with graphite electrodes, the anodic electrode was inserted in a small piece of coyonoztle (Opuntia imbricata). MFC was operated in fedbatch mode during a period of 60 days, using glucose as carbon source glucose (3 g/L), three external loads were used (5.5 KΩ, 1 KΩ and 560 Ω). Maximum voltage, current and power density was obtained for the case of 1 KΩ: 205.88 mV, 139.33 A y 3.43 W.cm², respectably.

MATERIALS AND METHODS

Experimental set-up: The two-chamber MFC were constructed with two plastic bottles with a volume of 1000 mL each. Plastic bottles were perforated and attached an inner screw thread joint to each bottle. An external thread joint (4 in length) was use to constructed the agar salt bridge (agar 3%, KCl 100 Mm). Graphite rods (Ø 0.8 cm×4 cm length, surface area 10.55 cm²) were...
used as electrodes for both chambers and connected using coated copper wire 22 gauge and sealed with silicon.

**Support media:** Opuntia imbricata was collected from the surrounding of city of Saltillo, Coahuila, Mexico. Small cylindrical pieces of dried stems of *Opuntia imbricata* were cut [(6 cm length, 5 cm O.D. 2.5 cm I.D.) ±0.2 cm] and meticulous rinsed and washed with fresh water, subsequently it was washed with distilled water to remove any present matter, and finally it was subject to sterilization for a period of 1 h. An electrode was inserted in the support media.

**Biofilm growth:** Support media with electrode was placed in a 1 L container and 250 mL of anaerobic sludge obtained from a UASB reactor and 500 mL of medium containing glucose as the electron donor (3 g/L = 850 mg COD/L), in a mineral medium Na₃H₂PO₄·2H₂O (0.53 g/L), NH₄Cl (0.3 g/L), MgCl₂·2H₂O (0.1 g/L), CoCl₂·6H₂O (0.08 g/L), NiCl₂·6H₂O (0.02 g/L) and ZnCl₂ (0.001 g/L) was added. After 4 weeks assembly of support media and electrode was withdraw from the container and placed in the MFC. Cathodic solution KMnO₄ (20 mm) was used. Evaluation of MFC was conducted at room temperature (18-23°C).

**Data acquisition and measurements:** Voltage (V) was measured every 60 min with a digital multimeter with data logging capacity (FLUKE 287), current (I) was calculated according to Ohm’s Law, \( I = \frac{V}{R} \), where \( V \) is the measured voltage and \( R \) is the external resistance (5.8, 1 and 0.560 KΩ). Maximum power density was determined as described by Lyon et al. (2010), \( P = VI/A \); where \( A \) (m²) is the surface area of the electrode. The glucose overall removal as COD was determined as describe by Rodriguez-Martinez et al. (2005).

**RESULTS AND DISCUSSION**

**Voltage and current generation in MFC:** MFC was operated under fedbatch and an external resistance was always present for the entire duration of the experiment. At the beginning for the first 672 h the external resistance was 5.8 KΩ. During this time the voltage generated by the MFC never passed the 100 mV with a maximum voltage of 94.1 mV as showed in Fig. 1. After 200 h MFC potential started to increase reaching its maximum at 490. After MFC was fed at 462 h, it reached its maximum, and a drop in the MFC potential was observed. This could be a possible shift in microbial community as suggested by Reguera et al. (2006), they found that many cells were not actively contributing to current production and current production was relatively low to the amount of biomass on the anode (Reguera et al., 2006; Biffinger et al., 2007; Marcus et al., 2007), and also it has been mentioned that microbial population is more diverse when MFC is fed by carbohydrates, this probably been a factor in the drop in the potential by presence of more non-electrogenic microorganisms in MFC (Jang et al., 2010).

As expected, when external resistance was changed to 1 KΩ an increase in the anodic potential was observed (Fig. 1, 2 and 3) this could be explained due to stimulation of the reoxidation of any electron mediators that may be present in the system, hence facilitating the electron transfer from bacteria to such mediators. In general, the use of smaller external resistance caused an increase in electron flow (Marcus et al., 2007; Cheng et al., 2008; Lyon et al., 2010).
The maximum power density reported here is modest compared with MFC using a proton exchange membranes, due to higher internal resistance of the salt bridge system as describe by Min et al. (2005), where comparison between MFC with two different selective membranes were conducted (proton exchange membrane vs. salt bridge), the low power output by the MFC with a salt bridge is directly attributed to the high internal resistance (1992±50 Ω) compared to the low internal resistance by the MFC with proton exchange membrane (1286±1 Ω). However, early reports by our research group, suggests the results obtained in this study are found to be very promising for a salt bridge two-chamber MFC (Guerrero-Rangel et al., 2010). In addition porous structure of Opuntia imbricata could possibly contributed in higher current density compared to early reports by our research group (Guerrero-Rangel et al., 2010) due to the fact that porosity and roughness of Opuntia imbricata’s structure enhanced biofilm formation, also the existence of a heterogeneous structure creates open channels that allowed the bulk fluid to penetrate to variable depths within biofilms, resulting in a better substrate access (Hu et al., 2005; Liu et al., 2010).

CONCLUSION

In order to be consider a practical use giving power output over long periods of time MFC will have to be converted to continuous flow and employing substrates such as wastewater and variety of organic pollutants. The use of expensive catalysts for the cathode must also be avoided, being oxygen from free air the ideal catalyst. In such systems, substrate and other nutrients will be continuously supplied to the bacteria and furthermore, there will be no waste product accumulation as these will be constantly driven out of the system. Designing MFC to operate in a continuous mode is a challenge that will have to be addressed according to the type of MFC under consideration.

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REFERENCES


