MICROBIAL FUEL CELL (MFC) TECHNOLOGY FOR HOUSEHOLD WASTE REDUCTION AND BIO-ENERGY PRODUCTION

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ABSTRACT

MFC is a bioreactor, extracts chemical energy from organic compounds, directly as electrical energy, through microbial degradation under anaerobic conditions. The main objective of the current study is to compare the degradation ability and corresponding electric potential development from different household substrates using lab scale MFC. 50hr batch experiments were conducted with household organic rich substrates like coconut water, rice starch and milk. Different concentrations of KMnO₄ were used as oxidizing agent in the cathode chamber. A voltage of about 300 to 700mV was produced from 125ml of substrates seeded with cow dung. Coconut water and starch produced electric potential with the support of oxidizing agent KMnO₄, whereas the potential produced by milk found to be independent of the KMnO₄ concentration. The maximum electric potential developed was 762mV from coconut water at 1500mg/l KMnO₄ with a COD reduction of 22%.

KEYWORDS

Bio-Energy, Household waste organic substrates, Microbial degradation, Electric potential, Oxidizing agent

1. INTRODUCTION

Industrialization and growth of population have resulted in severely increased demand of fossil fuels for energy. The increase of energy cost, depletion of non-renewable energy sources like fossil fuels and intolerable changes in the climate have considerably motivated scientists to work on finding new alternative energy production approaches. The other by-product of development is wastewater, produced from different sectors [1]. Hence Green energy is the need of the day. In the field of wastewater treatment, the conventional approach is to remove the pollutants or contaminants from wastewater and discharge the effluent to any water bodies. Here the challenges are of multiphase. The partially treated effluent from wastewater treatment plants affects the quality of water reserves, the separated contaminants create disposal problems and along with that treatment plants consume huge amount of energy. Microbial Fuel Cell (MFC) Technology is an approach in which, wastewater is considered as a resource, where wastewater is cleaned for re-use and energy is produced from the contaminants with a lower carbon footprint. Biodegradable organic-rich wastes from domestic, industrial and agricultural sectors are the ideal candidates as green energy sources for electrical production, since energy is biologically extracted from them and the wastes are treated concurrently.
MFC is a bioreactor that extracts chemical energy in the chemical bonds of organic compounds directly as electrical energy through catalytic reactions of microorganisms under anaerobic conditions. MFCs produce electricity from organic waste directly, without the need for methane collection, its conversion to heat by combustion and then to electricity. The bio-conversion in MFCs can occur at low substrate concentration levels and at temperatures below 20°C, where anaerobic digestion generally fails due to low reaction rates and high solubility of the methane produced. Power generation of MFC depends on many factors including type of membrane, catalyst, substrate, configuration, temperature etc [2].

Typically, MFCs, as it is shown in Figure 1, consist of anode and cathode chambers, separated by a proton exchange membrane (PEM). Organic matter or biomass is oxidized under anaerobic condition at the anode producing CO\(_2\), protons as well as electrons. Microorganisms here fulfil the role of biocatalysts. The protons are diffused to the cathode chamber through the PEM or CEM (Cation Exchange Membrane). Electrons produced by the bacteria from these substrates are transferred to the electrode (anode) and flow to the cathode (positive terminal) linked by a conductive material containing a resistor or device (like LED Bulb). Oxygen has usually been a final electron acceptor in the cathode due to its convenience, strong oxidation potential, and not being a chemical waste product. The electrons that reach the cathode through external circuit combine with protons and oxygen and the resulting product is water [3, 4, 10].

![Figure 1. Schematic Sketch of Microbial Fuel Cell [5]](image)

The bio-chemical reactions taking place in anode and cathode cells of MFC to create energy can be explained by Eq. (1).

**Anode reaction**: \[ \text{C}_6\text{H}_{12}\text{O}_6 + 6 \text{ H}_2\text{O} \rightarrow 6 \text{ CO}_2 + 24 \text{ H}^+ + 24 \text{ e}^- \]

**Cathode reaction**: \[
\begin{align*}
24 \text{ H}^+ + 24\text{ e}^- + 6\text{ O}_2 &\rightarrow 12 \text{ H}_2\text{O} \\
\text{C}_6\text{H}_{12}\text{O}_6 + 6 \text{ O}_2 &\rightarrow 6 \text{ CO}_2 + 6 \text{ H}_2\text{O} + \text{Electrical energy} \quad \ldots \ (1)
\end{align*}
\]

Electrical energy theoretically approaching – 2840 kJ/mol. [6]

Oxygen is having very slow reduction kinetics on graphite; therefore, it is one of the restricting agents in MFCs [7]. To alleviate this challenge, Potassium Ferri-cyanide (K\(_3\)[Fe(CN)\(_6\)]) is the commonly used oxidizing agent in most of the researches [4,8,9]. The two chambered MFC, using artificial sucrose wastewater as the substrate and hydrogen-producing mixed bacteria as the anodic inoculums, aerated catholyte and potassium ferricyanide catholyte were operated separately to differentiate the power generation. The paper reported that the reactor with potassium ferricyanide catholyte produced higher power. The effect of potassium ferricyanide concentrations in catholyte (pH 7.0, 100 mM phosphate buffer solution) on electricity generation
characteristics was reported that the maximum resultant MFC output power density of 181.48 mW/m² was produced using a potassium ferricyanide concentration of 0.1 M in the catholyte.

In a two chambered mediator less MFC using glucose as substrate and three different electron acceptors potassium permanganate (KMnO₄), potassium ferricyanide (K₃[Fe(CN)₆]) and potassium dichromate (K₂Cr₂O₇) were compared. The salt bridge connected MFC reported to produce a maximum potential of 1.04 V with KMnO₄. Performance of MFC with K₃[Fe(CN)₆] and K₂Cr₂O₇ were much lower with a maximum potential of 0.71 V and 0.56 V respectively.[11] The present study was conducted on a recreated version of MFC using locally available substrates. Worldwide, several studies were reported on analyzing MFC using synthetic organic wastes. But this study was conducted with a MFC system using natural substrates like coconut water, starch drained from boiled rice and milk solution (50% dilution) as substrates, which are available and generally thrown away from 90% of households and restaurants of Kerala. Being a poisonous chemical, Potassium Ferri-cyanide (K₃[Fe(CN)₆]) was replaced with KMnO₄ as the oxidizing agent in cathode chamber at different concentrations.

2. EXPERIMENTAL INVESTIGATIONS

2.1 Materials

Acrylic is a useful, clear plastic that resembles glass, but stronger than glass and resistant against chemical attack. A lab scale, two chambered MFC model was designed having a capacity of 125ml each. The connecting side of each chamber was provided with 30mm diameter hole, where the cation exchange membrane separates the chambers. The cation exchange membrane, CMI-7000 (Membranes International Inc., NJ, USA) was used for the internal conductivity between chambers. Graphite electrodes connected with light gauge wires were used for the external circuit of MFC. The anode and cathode chambers were connected by nuts and bolts. To prevent leakage between chambers rubber gaskets with 30mm hole at the centre were used and sealed with M-seal.

Coconut water, boiled rice drain water (starch) and milk solution were used as organic substrate in the anode chamber. Different concentrations of KMnO₄ were used in the cathode chamber. A Digital Multimeter was used for the measurement of electrical potential between electrodes. Initial and then final COD of the substrate samples were evaluated to find the treatment efficiency.
2.1 Methodology

Experimental evaluation was done in the microbial fuel cell (MFC) lab model as shown in Figure 2. The experimentation was done in three stages using three different substrates: Coconut water, boiled rice drain water (starch) and milk residue. In the first stage, coconut water was used as the substrate for biodegradation and hourly electric potential generation was monitored for 50 Hours as a batch reactor. Cathodic chamber was loaded with 500mg/l, 1000mg/l, 1500mg/l and 2000mg/l of KMnO₄ and on controlled system without KMnO₄. Inoculum for seeding was prepared, by adding 245 ml of coconut water with 5ml cow dung, one day before setting up of the cell. While loading the MFC, 115ml of fresh coconut water was seeded with 10ml of the prepared inoculum. The experiment was carried out without any control over temperature or pH. The same procedure was repeated using boiled rice drain water (starch) and milk solution as substrate with 50Hrs batches. Developments of Electrical potential, with respect to different concentrations of KMnO₄, were evaluated. Hourly Voltage fluctuations were measured using Digital Multi-meter. The measurements were plotted to compare the effect of concentration of oxidising agentKMnO₄ in the cathodic chamber on Electric potential developed (in mV). The values were compared with a control batch in which, the aerobic condition on cathode chamber was dependent only on natural surface aeration.

3. RESULTS AND DISCUSSIONS

3.1 MFC using Coconut water as substrate

The variation in electric potential developed between anode and cathode, using coconut water as substrate, with respect to time is represented in figure 3. The change in concentration of KMnO₄ showed significant effect in the potential developed. Optimum concentration of KMnO₄ is 1500mg/l producing a maximum potential of 762mV (half the potential of common battery). Potential development except in 1000mg/l, showed a similar pattern of steep increase at initial time period either ended at a steady state or decrease after a steady state, within 50hrs.

![Electric Potential vs Time](image)

Figure 3. Hourly variation in Electric Potential developed in MFC using Coconut water as substrate and different concentrations of KMnO₄ as cathode medium
The figure 3 showed that there was significant effect on the availability of proton accepter in cathode chamber, in the development of potential. Since biodegradation rate of substrate in the anode chamber produce electrons, higher concentration of KMnO₄ will not influence the development of potential.

### 3.2 MFC using boiled rice drain water (starch)as substrate

The variation in electric potential developed between anode and cathode, using boiled rice drain water (starch) as substrate, with respect to time is represented in figure 4. The change in concentration of KMnO₄ showed significant effect in the potential developed. Optimum concentration of KMnO₄ was found to be 1000mg/l producing a maximum potential of 335mV. Potential development showed a similar pattern of steep increase at the initial time period and ended at a steady state of around 250mV, within 50hrs.

![Electric Potential vs Time](image)

**Figure 4.** Hourly variation in Electric Potential developed in MFC using starch as substrate and different concentrations of KMnO₄ as cathode medium

The figure 4 clearly indicated the effect on the availability of proton accepter in cathode chamber, in the development of potential. Similar to MFC using coconut water as substrate, high concentration of KMnO₄ was not influenced in the development of electric potential. The lower potential development in the system compared to coconut water was due to the complexity and suspended nature of organic substances in the starch solution. But it gives a steady potential for a very long period, slow but steady biodegradation.

### 3.3 MFC using milk solution as substrate

The variation in electric potential developed, using milk solution (50% diluted milk) as substrate, with respect to time is represented in figure 5. At a KMnO₄ concentration of 1000mg/l, a maximum potential of 389 mV was observed. But a steady increase in Potential was achieved in the absence of KMnO₄ to the maximum of 429mV.
MFC using milk showed a similar pattern of steep increase at initial time period followed by a steep decrease in presence of KMnO₄ within 50hrs. In the absence of KMnO₄, a steady but slow increase and a similar decrease was observed. The physical examination showed formation of settled floc in the anode chamber. Hence it revealed the effect of curding in the development of electric potential rather than biodegradation in presence of KMnO₄. Electrons produced by biodegradation from the settling floc cannot be effectively transferred to the electrodes and hence showed a steep decrease in electric potential.

### 3.4 COD removal Efficiency in MFC

Table 1 showed the COD reduction efficiencies of different substrates in MFC compared to the natural anaerobic reduction of COD of the same substrates. Initial COD of substrates were found to be 48000mg/l for coconut water, 31000mg/l for starch and 50000mg/l for Milk solution. Approximately 50% increased rate of biodegradation can be observed in MFC compared to natural anaerobic biodegradation in all substrates. This is due to the favourable condition (balance of pH) for microbial growth created in MFC by the transfer of excess H⁺ ions from anodic chamber to cathodic chamber through CEM. Starch shows the largest % reduction of
COD as it was the with the substrate lowest initial COD concentration. Since potential difference influence the migration rate of H$^+$ ions, slight changes are observed in the degradation efficiency of MFC with respect to concentration of KMnO$_4$. In milk solution as substrate similar to the highest electrode potential, highest COD removal efficiency also associated with 0mg/l of KMnO$_4$. This result also prove that the migration efficiency of H$^+$ ions, in MFC has a significant role in the rate of biodegradation.

4. CONCLUSIONS

The Microbial fuel cell developed using household waste organic substrates was found to be developing higher electric potential by degradation with the support of proton accepter, KMnO$_4$. A steady electric potential in the range of 700mV was developed using Coconut water as substrate with an optimised KMnO$_4$ concentration of 1500mg/l within 30hrs. The lower range of steady electrical potential development (250 – 300mV) was observed in MFC using starch and milk as substrate with the support of proton accepter, KMnO$_4$. This is due to the presence of complex organic substances in colloidal state and its slow biodegradability. The observation from this study can be concluded that:

- Increase in concentration of proton accepter/oxygen releasing substances, increase the ability to extract potential from MFC at lower concentration, but beyond some limit, the influence reduces. The optimum level of oxygen or oxidising agent depends on the substrate and its rate of biodegradation. In MFCs, the optimum concentrations of KMnO$_4$ as proton accepter were 1500mg/l and 1000mg/l for substrates coconut water and starch respectively.

- When milk was used as substrate, the presence of KMnO$_4$ as oxidising agent make curding, to reduce the potential (250 – 300mV) that is extracted from MFC. This was due to the change in phase of organic materials, from dissolved to suspension, which restrict biodegradability and the transfer of electrons developed during biodegradation to electrode. In the absence of KMnO$_4$, the organic components retain in dissolved form and produced comparatively higher steady potential in the range of 350 – 400mV.

- COD removal efficiency was approximately doubled irrespective of the KMnO$_4$ concentration in anodic chamber. Hence H$^+$ ion migration in MFC from anode to cathode chamber through CEM created a favourable condition for biodegradation.

The system needs further optimization study in terms of electrode material, its structure, membrane surface area, oxidising agent used in the cathode chamber and chamber design. The electric potential of MFC can be increased by efficient seeding with specific anaerobic bacteria type suitable to substrate. Other scope is the use of mediators to improve electron conductivity from bacteria to electrode.

ACKNOWLEDGEMENTS

The authors would like to thank Aroma Plastics, Kochi for their support for casting the reactor chambers as per required design and friends for the support in purchase of Cation Exchange Membrane from Membrane International, Ringwood, USA.
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