Dynamic evolution of anodic biofilm when maturing under different external resistive loads in microbial fuel cells. Electrochemical perspective

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**HIGHLIGHTS**

- Dynamic changes of biofilm matured under different external loads are investigated.
- Most dynamic changes are observed in the first 3 weeks of operation.
- External resistance lower than internal resistance improves maturing of the biofilm.
- Internal resistance is dependent on external resistance applied in maturing phase.
- External resistance irreversibly affects the biofilm properties and performance.

**GRAPHICAL ABSTRACT**

**ABSTRACT**

Appropriate inoculation and maturation may be crucial for shortening the startup time and maximising power output of Microbial Fuel Cells (MFCs), whilst ensuring stable operation. In this study we explore the relationship between electrochemical parameters of MFCs matured under different external resistance ($R_{\text{ext}}$) values (50 Ω - 10 kΩ) using non-synthetic fuel (human urine). Maturing the biofilm under the lower selected $R_{\text{ext}}$ results in improved power performance and lowest internal resistance ($R_{\text{int}}$), whereas using higher $R_{\text{ext}}$ results in increased ohmic losses and inferior performance. When the optimal load is applied to the MFCs following maturity, dependence of microbial activity on original $R_{\text{ext}}$ values does not change, suggesting an irreversible effect on the biofilm, within the timeframe of the reported experiments. Biofilm microarchitecture is affected by $R_{\text{ext}}$ and plays an important role in MFC efficiency. Presence of water channels, EPS and precipitated salts is distinctive for higher $R_{\text{ext}}$ and open circuit MFCs. Correlation analysis reveals that the biofilm changes most dynamically in the first 5 weeks of operation and that fixed $R_{\text{ext}}$ lefts an electrochemical effect on biofilm performance. Therefore, the initial conditions of the biofilm development can affect its long-term structure, properties and activity.

1. Introduction

Microbial Fuel Cell (MFC) technology uses electroactive bacteria to produce electricity through oxidation of organic matter. The technology has received increased attention over past decades [1]. The bioelectrochemical reactions take place in anodic and cathodic components of
the MFC have found many potential applications in the fields of wastewater treatment, electricity generation, biogas production, biosensors and bioelectrochemical synthesis [2–10]. Among various carbon sources that have been demonstrated as a fuel in MFCs human urine has proved to be a good substrate due to its high conductivity [11]. The ongoing development of the MFCs focuses on developing electrode materials, catalysts, membranes separating anodic and cathodic chambers, design and scale-up of the MFC-based systems [2,12–17].

Despite the broad interest in many engineering aspects of the MFCs, the most crucial role is played by the electroactive bacteria, which form the biofilm on the electrodes' surface and generate electrical power from their population-level metabolism. The biofilm is a complex matrix of microorganisms and extracellular compounds which is considered to be very stable, albeit possessing physiological adaptive mechanisms [18] many of which are expressed during the initial biofilm formation period.

Several research groups have previously reported on power performance and start up times when the electroactive community has been matured under different poised anode potentials. For example the anodic biofilm formed by Geobacter sulfurreducens gives highest power performance and lowest MFC start-up time, when matured under a potential range between 0 and 400 mV vs SHE (standard hydrogen electrode). This optimal potential range promoted the biofilm growth and corresponding power density of the MFCs [19]. The study reported by Aelterman et al. showed, that optimal biofilm growth and activity was obtained when the anode was poised at −200 mV vs Ag/AgCl electrode, although the original source of the bacterial inoculum was not mentioned [20]. More recently, Zhu et al. reported, that acclimating the biofilm with positive potentials may lead to the decay of the power overshoot phenomenon which leads to improved power performance [21].

Easier ways of controlling the potential of MFC electrodes is by applying an external load (Rext), which does not require any specialist equipment that could be limiting in particular for field applications. Comprehensive investigation on the effect of Rext on biofilm formation and activity has been reported by Zhang et al. [22]. The authors investigated the ohmic range of 10–1000 Ω and indicated that optimal Rext for their MFC setup was found to be 50 Ω, although biofilm matured under 10 Ω produced the highest current. The study also showed that Rext had an impact on the presence of extracellular polymeric substances (EPS) of the biofilm, and a more recent study showed that EPS plays a role in biofilm performance and in turn, power generation [23]. The influence of three different Rext values on biofilm activity (after the maturing phase) was also studied by Jung and Regan [24]. The authors focused on methane production and the inhibition of methanogenesis was found to occur in parallel with the highest power efficiency for MFCs fed with acetate and operating under lowest (150 Ω) Rext. Earlier studies also demonstrated the relationship of Rext applied during operation (after maturing phase) with performance of the MFCs in relation to the fuel supply and the best results were obtained when MFCs were operated under Rext closer to internal resistance (Rint) [25]. External resistance was also found to be a factor influencing diversity of the bacterial community [24,26,27].

Although significant work has been done to understand the interactions of Rext with the biofilm, very limited knowledge is available on dynamic evolution of biofilm subjected to various external loads. Since the biofilm forms both stable and adaptive structure, such knowledge is indispensable to develop appropriate strategies for inoculation and operation of MFCs. It is therefore important to determine, whether the conditions applied to the biofilm in the initial stage of development may leave a structural and electrochemical profile and irreversibly affect its performance thereafter.

The aim of this study was to determine the temporal and long-term effects of fixed and dynamically-changed external resistance on changes of biofilm parameters and resulting MFC performance in time. The results revealed the irreversible effects that the initial Rext causes to the biofilm, which may subsequently either induce or inhibit the power performance of the MFCs in long-term perspective. This is the first of two papers in series, where we have focused on analysis of electrochemical parameters. The second part of this study will focus on biological parameters of the biofilm.

2. Experimental

2.1. MFC construction and operation

The single chamber MFCs were built as described in detail in our previous work [28]. In brief, earthenware ceramic material was used both as the separator and housing for the anodic chamber (Fig. 1). The external side of the ceramic cylinder was supplied with carbon-painted cathode (carbon loading of 14.08 mgC cm\(^{-2}\)) and stainless steel wire mesh, acting as a current collector. The volume of an empty MFC anodic chamber was 11.4 mL. Carbon fiber veil (20 g m\(^{-2}\), PRF Composite Materials, Dorset, UK) was used as the anodes with total surface area of 252 cm\(^2\). A 3D-printed Nanocure™ RCP30-resin lid with inlet and outlet tubes was used as a front panel of the vertically positioned MFCs.

All of the MFCs were manufactured manually in the same manner and used to build an array of 21 electrically and fluidically isolated
MFCs. The array consisted of 7 sets of triplicate MFCs. Each triplicate was operating under different electrochemical conditions, controlled by the external load, connected to each individual MFC.

In the first stage of experiments, the external resistance ($R_{\text{ext}}$) was predefined for each triplicate, as shown on Fig. 1. The array consisted of MFCs connected to 50, 500, 1000, 5000 and 10000 $\Omega$, as well as the open circuit (OC) control. The electroactive biofilm was grown at the anodes using these predesigned $R_{\text{ext}}$ values for approximately 6 weeks. Activated sludge derived from the aerobic chamber of municipal wastewater treatment plant (Wessex Water, Saltford, UK) was used to inoculate the MFCs. After 24 h, activated sludge was removed from the MFCs and replaced with the fresh human urine as a fuel. The MFCs were operated in continuous flow conditions. The fuel was delivered to the MFC-array using multichannel peristaltic pump (Watson Marlow, USA) at a constant flow rate of 0.12 L d$^{-1}$. Between stages 1 and 2, the MFCs were fed with gradually increasing flow rates to determine the impact on power performance. Before the second stage, the flow rate was brought back to a constant value of 0.12 L d$^{-1}$.

The second stage of the experiment took place between 7 and 17 weeks of operation and was conducted in the same hydraulic conditions. In the second stage, the $R_{\text{ext}}$ values were continuously adjusted to the optimum (simulating MPPT). Each graph represents three individual MFC replicates.

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2.2. Dynamic electrochemical behaviour of the biofilm

Electrochemical behaviour of the biofilm was tracked over time through polarisation experiments and real time temporal power performance monitoring. Polarisation experiments were carried out during each week. In the second stage, the optimal $R_{\text{ext}}$ values determined from power curves were applied to each MFC to simulate maximal power point tracking (MPPT) conditions. The polarisation experiments were conducted using automated variable resistor system, known as the resistorstat [29]. Each polarisation run consisted of 1 $\Omega$–3.75 $\Omega$ resistance range. Each value was connected to the MFC for a period of 5 min. The power performance of the MFCs was monitored with 34972 A Data Acquisition unit (Agilent Technologies, USA) with the data logging sample rate set to 3 min. The current was calculated according to Ohm's law: $I = V/R$, where $V$ is the measured voltage in Volts (V) and $R$ is the value of the external resistance. The power output $P$ in Watts (W) was calculated using equation: $P = I \times V$.

2.3. Environmental electron scanning microscopy

The biofilm structure was investigated by field emission environmental scanning microscopy, coupled with Energy-dispersive X-ray spectroscopy (Philips XL-30). Samples of the biofilm were collected from the anode at the end of the experimental period and fixed with 4% glutaraldehyde in 0.1 M PBS buffer. Subsequently, the samples were rinsed with water and air-dried for 12 h at room temperature.
2.4. Data processing and statistical analysis

Experimental data were first processed using Microsoft Excel 2010 and further analysed and visualized using R GUI package (v. 3.4.2). The polarisation experiments data was processed and visualized using SciDAVis (v. D001) software. Investigating the dynamic effects of Rext on the biofilm properties and activity was conducted by local polynomial regression fitting, as well as by determining the Pearson’s correlation coefficients between Rint, Rext, OCV (open circuit voltage), power and current, extracted from polarisation experiments. The data was further visualized through the correlogram.

3. Results and discussion

All experiments were carried out using fresh human urine, collected on a daily basis as a fuel and delivered under constant flow rate. Therefore, the measured pH of fresh urine ranged from 6.15 to 6.29 and the average conductivity was equal to 11.76 ± 0.76 mS.

3.1. Electrochemical behaviour of biofilm

It is known, that the shape of a polarisation and power curves and the corresponding losses are greatly dependent on the activity of electroactive bacteria [32,33]. Data derived from the polarisation experiments were presented in the form of power curves, as shown in Fig. 2. The results reported in this study comprised the highest coverage of Rext within the operational range of MFCs when compared to previous studies available in the literature. Previous works used up to four Rext values below 10 kΩ. The diode effects of Rext values below 10 kΩ are susceptible to starvation (0.008 L d⁻¹), which led to a drop of power output when compared to other Rext. These data perhaps suggest that the 50Ω biofilm had developed mechanisms for more efficient conversion of nutrients into electricity when the feeding rate is sufficiently high, but underdeveloped the ability to accumulate store energy reserves in such form as EPS (Fig. 3) which could be used when the feeding rate is low. Flow rate is an important parameter that is bound to affect the biofilm as a whole. Previous reports [41] describe how the flow rate affects the biofilm growth rate and in turn power output. However, the biofilm structure and how this may be affected by flow rate, is an area that requires dedicated investigation.

After 5 weeks of operation (stage 2), the pre-defined resistor values were replaced by the optimal Rext values (derived from the polarisation data), adjusted to the optimum value at one-week intervals (simulating an MPPT approach). Changing the Rext of the MFCs to follow their electrochemical properties [40].

The differences in biofilm properties were also revealed during the flow rate experiment (Figure S2). The results indicate, that only the 50Ω-matured biofilm was capable of utilising nutrients at the highest flow/supply rate (0.888 L d⁻¹). In parallel, the 50Ω-biofilm was more susceptible to starvation (0.008 L d⁻¹), which led to a drop of power output when compared to other Rext. These data perhaps suggest that the 50Ω biofilm had developed mechanisms for more efficient conversion of nutrients into electricity when the feeding rate is sufficiently high, but underdeveloped the ability to accumulate store energy reserves in such form as EPS (Fig. 3) which could be used when the feeding rate is low. Flow rate is an important parameter that is bound to affect the biofilm as a whole. Previous reports [41] describe how the flow rate affects the biofilm growth rate and in turn power output. However, the biofilm structure and how this may be affected by flow rate, is an area that requires dedicated investigation.

After 5 weeks of operation (stage 2), the pre-defined resistor values were replaced by the optimal Rext values (derived from the polarisation data), adjusted to the optimum value at one-week intervals (simulating an MPPT approach). Changing the Rext of the MFCs to follow their optimal values (when Rext = Rinst) resulted in changes to the shape of MFC power curves. The ohmic losses previously observed in stage 1 were no longer visible. Such an interesting phenomenon suggests, that there was perhaps a change in biofilm behaviour that adjusted to the new electrochemical conditions. This could be explained by the increased oxidation of the carbon-energy substrate due to higher metabolic rates, since the new resistors connected to the MFCs had lower resistance, when compared to the stage 1. Higher metabolic rates are proportional to higher microbial growth rates and hence biomass density of the electroactive species. In a perfusable electrode system this maintains a fixed thickness biofilm, as the outer layers, which no longer have direct contact with the electrode, are constantly washed out. This results in a thinner biofilm and improves the diffusion of nutrients into the biofilm and conductivity (as indicated by Rinst – Fig. 3). A thinner biofilm will also result in a significant reduction in extracellular polymeric substances (EPS) as previously reported by Zhang et al. [22]. The present study, suggests that Rinst could dynamically change over time, as also alluded to by Winfield et al. [34].
3.2. Changes of power and internal resistance over time

To further investigate the impact of $R_{\text{ext}}$ on biofilm maturing, the dynamic changes in $R_{\text{int}}$ and power over time were also monitored (Fig. 3). In the first week of operation, the highest power performance was observed for 1 k$\Omega$-matured MFCs and reached 91.6 ± 14.0 $\mu$W, whilst the $R_{\text{int}}$ reached a value of 471.3 ± 247.5 $\Omega$. The lowest $R_{\text{int}}$ value was also observed for 1 k$\Omega$ and was equal to 205.4 $\Omega$ (derived from the polarisation experiment). Thus, in the first week, the use of 1 k$\Omega$ $R_{\text{ext}}$ created the most favourable conditions to obtain the lowest $R_{\text{int}}$.

During the first 3 weeks, the highest power performance was observed for 1 k$\Omega$ MFCs. In the following weeks the maximum power output was recorded for lower $R_{\text{ext}}$ values: 500 $\Omega$ MFCs in week 2 (81.3 ± 15.0 $\mu$W) and 50 $\Omega$ MFCs in week 3 (112.2 ± 23.1 $\mu$W). Similarly, the minimum $R_{\text{int}}$ values were observed for 1 k$\Omega$ and equal to 205.4 $\Omega$ (derived from the polarisation experiment). Thus, in the first week, the use of 1 k$\Omega$ $R_{\text{ext}}$ created the most favourable conditions to obtain the lowest $R_{\text{int}}$.

During the first 3 weeks of biofilm maturing, which appears to be the critical time required for the biofilm to develop. Furthermore, the highest performance of the MFCs during the entire experiment was also observed in the 3rd week of operation and was recorded for the MFCs matured under the lowest Ohmic value, 50 $\Omega$ (112.2 ± 23.2 $\mu$W). Following the 3rd week of operation, the increase of $R_{\text{int}}$ with increasing $R_{\text{ext}}$ was observed even in the 2nd stage of operation, where optimal $R_{\text{ext}}$ values were applied following the simulated MPPT method. The internal resistance showed similar (but negative) trends that could be observed for power performance, which was reflected by similar patterns developed by the MFCs through time (Fig. 3A–B). The correlation between those two factors was later confirmed by statistical analysis. The internal resistance may be affected by various parameters such as hydraulic and environmental conditions [25,42,43]. In this study the hydraulic and other operational conditions remained constant, even though the flow rate was relatively low for this type of MFC. Therefore, such dynamic changes of internal resistance, in particular in the early period of operation (stage 1) resulted from ongoing biofilm development. Such changes may affect the conductivity of the biofilm, which is directly correlated with the current density by reducing the resistance of the biofilm.
electron flow and lowering the activation energy required for electron transfer between biofilm and the anode [36].

Interestingly, at the end of the 1st stage, a local power optimum was observed for 2.5 kΩ and became even more distinctive in the 5th week of operation. The 2.5 kΩ MFCs reached 93.0 ± 7.1 μW, which corresponds to 89.8% of the performance observed for 50 Ω MFCs. This state was also observed in the later stage of the biofilm growth, when the optimal $R_{\text{ext}}$ resistors were applied to the MFCs (stage 2). Local polynomial regression fitting (Fig. 3A and 3. B) revealed that this local optimum could be found between $R_{\text{ext}}$ of 0.5 and 2.5 kΩ (with best performing MFCs observed for 1.0 kΩ) and was maintained until 15th-16th week of operation. At the same time 50 Ω MFCs, which appeared to show the highest power performance in stage 1, underperformed as compared to MFCs operating under 0.5–2.5 kΩ - matured MFCs. This can be explained by the occurrence of electroosmotic drag, which is related to the power output and could lead to faster accumulation of salts at the cathode observed in the best performing (50 Ω) MFCs [44,45]. In fact, the highest amount of salt deposits at the cathode surface were observed for 50 Ω MFCs, while the lowest for OC control and also 500–5000 Ω MFCs, indicating the interdependence between electroosmotic drag and MFC performance (Figure S1). The cathode performance adds another dynamic factor to the complexity of the microbial fuel cells.

3.3. Dynamic electrochemical profile - simultaneous effect of $R_{\text{ext}}$ and $R_{\text{int}}$ resistance on MFC performance

Data shown in Fig. 4 display a particular type of electrochemical profile – the pattern developed throughout time by all biofilm communities, reflecting dependence of the MFC electrochemical parameters on controlled (external) and developed (internal) resistance. The results
show that the highest power efficiency of the MFCs was achieved for the biofilm which developed an internal resistance lower than 300 Ω, and such low $R_{\text{int}}$ was only present when the biofilm was matured under $R_{\text{ext}}$ between 50 and 1000 Ω. Another local optimum was found for the biofilm matured under $R_{\text{ext}}$ between 1000 and 2500 Ω, which developed an $R_{\text{int}}$ between 407.2 and 701.8 Ω. Notably, biofilms developed under lower $R_{\text{ext}}$ values produced the highest current and the lowest OCV, while the biofilms matured under higher values of $R_{\text{ext}}$ showed lower current and higher OCV. These results could be explained by the fact that the biofilm subjected to the effects of the low external resistance (higher anode potential) develops a biofilm with a different composition of electroactive species, as shown by previous studies for both biofilm [38–40] and planktonic communities [38]. It was shown that the strategy of maturing the biofilm under $R_{\text{ext}}$ values lower than the lowest observed $R_{\text{int}}$ may be beneficial in the long term and this is (to the best of the authors’ knowledge) the first report where these phenomena have been demonstrated. Data reported in previous research focused on maturing the biofilm in $R_{\text{ext}} > R_{\text{int}}$ conditions [22,24–27]. Furthermore, when comparing profiles obtained in two stages of experiment, it can be concluded that the biofilm maintained its electrochemical properties even though the environment was dynamically changing over time. The obtained profiles were similar, including the general decreasing trend in performance, current and OCV, probably due to cathodic salt accumulation, as previously described. These findings are crucial in defining the appropriate inoculation strategy and protocols for assessing and predicting the MFC performance. The internal resistance proved to be a more effective parameter to determine the best performing MFCs than OCV. Nevertheless, reaching high OCV suggests that system may be better balanced and more effective in long term operation, when dynamic changes in cathode performance may occur. The data also shows, that reaching higher potential within MFC environment is not as important as acquiring the biofilm with a desirable structure and community composition as reflected by $R_{\text{int}}$, which allows the MFC to reach a higher current production rate and density. This finding suggests that the electrochemical biofilm properties developed over time, are as important as the composition of the electroactive consortia, which also changes dynamically over time [46–48].

3.4. Biofilm microarchitecture

The biofilm structure and composition was assessed at the end of experiment (after phase 2 - switching $R_{\text{ext}}$ to optimal values) using environmental scanning electron microscopy, as shown on Fig. 5. The most notable changes were in the form of inorganic precipitates. The MFCs matured under lower $R_{\text{ext}}$ (50, 500, 1000 Ω) showed no or negligible amounts of such precipitates. Maturing the biofilm under 2.5 kΩ and above resulted in increased amounts of graupel-shaped crystals embedded into the biofilm structure, comprising mainly Na, Cl, Mg, P and Ca, as determined by energy-dispersive X-ray spectroscopy. These salts were present in the highest amounts under open circuit conditions and in all reported cases, adjacent to neighbouring bacterial cells. Such finding may suggest a tendency from the anodic community to accumulate or induce formation of precipitates at very low or no current flow (see Fig. 5). This phenomenon could be one of the factors that contributed to the development of higher $R_{\text{int}}$ over time for MFCs that had matured under higher $R_{\text{ext}}$, since the crystals are considered to be non-conductive and may have affected the resistivity of the biofilm.

The differences in biofilm microarchitecture were also most notable when comparing high $R_{\text{ext}}$ (5 kΩ, 10 kΩ and OC) with low $R_{\text{ext}}$ MFCs. Images acquired for high $R_{\text{ext}}$ anodic biofilm and OC control, revealed development of looser structures, rich in EPS and with visibly larger water channels in comparison to the low $R_{\text{ext}}$ anodic biofilms. In contrast, the lowest $R_{\text{ext}}$ (50 Ω) MFC developed a dense and uniform biofilm structure with little EPS content. Zhang et al. [22], quantified the EPS content and reported its inverse relationship with external resistance, which is not in line with the findings of the current study. In that previous study, the results were normalised to the surface area of the carbon cloth electrode (as opposed to the biomass), in addition to using a different $R_{\text{ext}}$ range (10–1000 Ω).

In addition to the influence of $R_{\text{ext}}$ on the anodic community composition, as previously reported [24,26,27,49], the current study suggests that the biofilm microarchitecture was affected by $R_{\text{ext}}$ and played an important role in the evolution of the electrochemical MFC parameters over time, reaching a desirable power efficiency. The observed differences between the lower and higher ranges of $R_{\text{ext}}$, as well as the OC control, indicate that beyond a point, the biofilm was irreversibly

![ESEM images showing changes in biofilm three-dimensional architecture, composition and occurrence of inorganic precipitates (bright spots), when matured under different external loads.](image-url)
affected by its initial maturing conditions within the timeframe of experimental period.

3.5. Quantitative analysis of dynamic changes in biofilm behaviour

To investigate the dynamics of changes recorded for electrochemical parameters of the biofilm, correlation analysis has been conducted, as shown on Fig. 6. In the first stage of biofilm maturing, when fixed $R_{\text{ext}}$ values were applied, positive correlation ($R = 0.80$) was only found for current profiles recorded in the first two weeks, which we consider as a startup period of the MFCs. The correlation coefficient gradually decreased afterwards, confirming that most dynamic changes took place during the first 5 weeks of biofilm maturing. A similar pattern was observed for the power output, while the OCV profiles reached the highest correlation coefficients ($0.82$–$0.99$) among all of the parameters, starting from the 2nd week of biofilm maturing. In contrast, following the 2nd week a positive correlation was observed for $R_{\text{int}}$ ($0.35$–$0.76$) and was decreasing with the increase of time; i.e. two neighbouring $R_{\text{int}}$ profiles (variables) were showing the highest correlation. These results suggest, that internal resistance was smoothly changing from one state to another due to the development of the biofilm growth. It can be also concluded, that current was the most dynamic biofilm parameter, while the OCV was the most stable one. Therefore, the data suggest that the power output of the MFCs was rather not affected by the redox potential of bacterial enzymes and resulting electrode overpotentials. This statement is supported by finding a positive correlation between power and current ($0.51$–$0.93$) excluding week 2) and negative correlation with $R_{\text{int}}$ ($-0.38$ to $-0.91$), excluding week 9), while not even a weak correlation was found between OCV and power output. Interestingly, the OCV was very well correlated with the $R_{\text{int}}$ during the first weeks of operation reaching R values of $0.77$–$0.86$ following 2 weeks of biofilm maturing. Thus, $R_{\text{int}}$ was the main factor determining the potential of the MFCs and reaching high OCV values were not determining for the power performance.
When simulated MPPT method was used to control the power performance of the MFCs in stage 2 the biofilm had readjusted its metabolism to the new conditions. Since during this period the $R_{\text{ext}}$ was varying to fit the $R_{\text{int}}$, the dynamics of the changes in electrochemical biofilm properties have been partially inhibited. As a result, significant correlation was recorded for current throughout the second stage and reached between 0.40 and 0.89. The highest correlation coefficients were observed for the neighbouring time points and decreased with time, which suggest smooth (less dynamic) evolution of the biofilm from one state to another. In contrast, no significant correlation was observed when the changes of $R_{\text{int}}$ were followed in time (stage 2), while significant correlation for this parameter was observed in stage 1. This suggests that when the simulated MPPT procedure was applied, temporal and chaotic changes in $R_{\text{int}}$ occurred. Those changes which may have resulted from dynamic biofilm adaptation mechanisms coupled with ongoing changes in cathode performance. However, they were not reflected by the dynamic behaviour reported for current. Although strong negative correlation values were reported for $R_{\text{int}}$ and power in that period, power and current were also negatively correlated with original (stage 1) $R_{\text{ext}}$ where weak to moderate correlation was proved $R_{\text{int}}$ and power output. The most dynamic changes in electrochemical biofilm properties were observed in stage 2 during metabolic transition, when strong negative correlation for current was reported. Therefore, previous correlation coefficients were reported for 3rd week of operation, which we believe is the most significant period required to develop healthy and well-performing electroactive biofilm in MFC. Although MPPT method proved to be an efficient way of reducing the startup time and improving the performance of the MFCs, its application requires dedicated electronic circuit which a dedicated MPPT controller. The performance of a pilot-scale continuous flow microbial electrolysis cell fed winery wastewater, Appl. Microbiol. Biotechnol. 89 (2011) 2053–2063, https://doi.org/10.1007/s00253-011-3130-9.


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