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1 **Effect of Anode Polarization on Biofilm Formation**  
2 **and Electron Transfer in *Shewanella***  
3 ***oneidensis*/Graphite Felt Microbial Fuel Cells**

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8 **KEYWORDS.** Microbial Fuel Cell, *Shewanella oneidensis*, biofilm, bacteria/carbon  
9 interface, polarization.

10 **ABSTRACT.** In microbial fuel cell, electricity generation is assumed by bacterial  
11 degradation of low-grade organics generating electrons that are transferred to an electrode.  
12 The nature and efficiency of the electron transfer from the bacteria to the electrodes are  
13 determined by several chemical, physical and biological parameters. In particular, the  
14 application of a specific potential at the bioanode has been shown to stimulate the  
15 formation of an electro-active biofilm but the underlying mechanisms remain poorly  
16 understood. In this context, we have here studied the effect of an applied potential on the  
17 formation and electroactivity of biofilms established by *Shewanella oneidensis* bacteria on  
18 graphite felt electrodes in single- and double-chamber reactor configurations, in oxic  
19 conditions. Using amperometry, cyclic voltammetry and OCP/Power/Polarization curves  
20 techniques, we showed that a potential ranging between -0.3 V and +0.5 V (vs.  
21 Ag/AgCl/KCl sat.) and oppositely applied to a couple of electrodes leads to different  
22 electrochemical behaviors, anodic currents and biofilm architectures. In particular, when  
23 the bacteria were confined in the anodic compartment of a double-chamber cell, a negative  
24 applied potential (-0.3 V) at the bioanode favors a mediated electron transfer correlated  
25 with the progressive formation of a biofilm filling the felt porosity and bridging the  
26 graphite fibers. In contrast, a positive applied potential (+0.3 V) at the bioanode stimulates  
27 a direct electron transfer resulting in the fast-bacterial colonization of the fibers only. These

1 results provide fruitful insights for the understanding of the complex bacteria-electrode  
2 interplay in microbial fuel cells.

3

4

## INTRODUCTION

Microbial fuel cell (MFC) represents a promising energetical solution allowing the production of electricity from a diversity of organic substrates through direct oxidation by electrochemical active microorganisms under ambient conditions <sup>1</sup>. The power density output of an MFC is far lower than that of chemical fuel cell because the latter uses relatively clean energy sources such as hydrogen or methanol without implying other biological system, while MFCs typically use low-grade organics, such as domestic or industrial wastes. However, in return, using MFCs for waste treatment may significantly save energy <sup>2</sup>.

Many parameters influence the performance of MFCs. Besides the electrode materials and the used micro-organisms, there is accumulating evidence that the potential applied at the bioanode impacts on its colonization and therefore the MFC electrochemical performance <sup>3-8</sup>. However the intensity and the sign of the potential is often a matter of discussion <sup>9,10</sup> and highly dependent on the micro-organisms which can be a bacterium or a consortium artificially-build or sampled from real environment <sup>4,11</sup>, such as domestic or industrial wastewaters <sup>12</sup>, garden composts <sup>13</sup> or marine soils <sup>14</sup>. As an example, for *Geobacter sulfurreducens* positive applied potential (ca. + 0.26 V vs. SCE, between 0 to + 0.4 V vs. Ag/AgCl <sup>15</sup> or + 0.51 V vs Ag/AgCl <sup>16</sup> depending on the quoted study) was necessary. For mixed cultures and real inoculum, the behavior is often more complex, due to the multiplicity of bacterium strains, the synergic behavior of these strains and/or the domination of a specific strain. Moreover, the origin of inoculum influences its composition, and therefore its electrochemical response <sup>17</sup>.

Most of the time, the potential is empirically defined and, to date, no general law exists to predict the “ideal potential of polarization” with efficiency and accuracy. However, the use of a potential higher than the standard potential of the electron donor that is degraded is necessary <sup>18</sup>. Note that potential higher than the equilibrium potential favors electron deficiency at electrode surface, then encouraging oxidation reaction by bacteria <sup>19</sup>. In most cases, the polarization as well as the measurements are carried out in strict anoxic condition <sup>20,21</sup> to limit the competition with oxygen reaction for electron collection and to ensure the viability of anoxic strains such as *G. sulfurreducens*.

1 In this context, the aim of the present work is to study whether the potential applied at a  
2 bioelectrode constituted of *S. oneidensis* bacteria in contact with a graphite felt (GF) in oxic  
3 conditions affects or not the MFC performances. Two different MFC configurations were  
4 employed, single- and dual-chamber MFC configuration<sup>21,22</sup>, in order to apply opposite  
5 polarization at the two graphite felt electrodes considered as a bioanode<sup>23</sup> and a cathode<sup>24–</sup>  
6<sup>27</sup>. The applied potential was varied from - 0.3 V to + 0.5 V (vs. Ag/AgCl/KCl sat.) at the  
7 bioanode and its influence on biofilm formation and MFC electrochemical performances  
8 was studied. Electrochemical activity was measured by monitoring electrodes open-circuit  
9 potential, current density, MFC open-circuit voltage, and MFC polarization curves  
10 (maximal current and power)<sup>28</sup>. Moreover, the characterization of the biofilm organization  
11 after several days of functioning permitted to establish correlations between the  
12 electrochemical response and the colonization progress that was found to be sensitive to the  
13 sign and intensity of the applied potential.

## 14 15 MATERIALS AND METHODS

16  
17 All chemicals are bioreagent grad, provided by Sigma-Aldrich France and used without  
18 further purification. The graphite felt is gratefully delivered by Morgan Carbon company  
19 (Luxembourg)<sup>29</sup>. *S. oneidensis* CRBIP17.141 bacterial strain is prepared and delivered by  
20 Biological Resource Center of the Pasteur Institute (France).

21 **Inoculum.** *S. oneidensis* inoculum is prepared following two steps of pre-growth and  
22 growth. A fraction of the strain, conserved at -80°C, is inoculated and pre-cultivated into a  
23 Luria-Bertani Broth medium stirred at 150 rpm for 24 h at 30°C. Then, 1 mL of the pre-  
24 cultivated bacteria is transferred into a 50 mL MR1 growth-medium (see SI for complete  
25 description) with 30 mM sodium lactate and sodium fumarate as electron donor (carbon  
26 source) and electron acceptor, respectively. Bacterial growth is carried out for 18 h in oxic  
27 condition until reaching an optical density (at 600 nm, OD<sub>600</sub>) corresponding to the last  
28 quarter of the log-phase (OD<sub>600</sub> = 1.7) of the bacterial growth. The final inoculum is  
29 prepared by transferring the obtained bacterial pellet into fresh MR1 medium supplemented  
30 with 30 mM sodium lactate and free of sodium fumarate. The inoculum is stored at 4 °C in

1 sterile oxic condition for 10 min before being used as electrolyte for electrochemical  
2 reactor.

3 **Single- and dual-chamber reactor set-up and polarization experiment.** In both single-  
4 and dual-chamber reactors, GF was used as anode ( $1 \text{ cm}^3$ ) and cathode ( $2 \text{ cm}^3$ , to ensure no  
5 current limitation due to the cathodic electro-active surface). A three electrodes  
6 configuration was employed for electrodes polarization and for electrochemical  
7 characterization using Ag/AgCl/KCl sat. reference electrode.

8 The single-chamber configuration was composed of a 30 mL sealed vessel. The GF anode  
9 and cathode were used in 20 mL of the *S. oneidensis* inoculum diluted at 0.7 in OD<sub>600</sub>,  
10 corresponding to  $8 \cdot 10^8 \text{ cfu.mL}^{-1}$ . For all experiment, *S. oneidensis* inoculum corresponds to  
11 the reactor electrolyte composed of bacteria dispersed in a MR1 medium (growth medium,  
12 as previously described) supplemented with 30 mM lactate and free from fumarate,  
13 buffered at pH 7.

14 For the dual-chamber configuration, two compartments were separated by an  
15 ultrafiltration membrane (pore size:  $0.2 \mu\text{m}$ ). The anodic side was filled with a bacteria  
16 solution (as previously described) and the catholyte consisted in a solution of 10 mM  
17  $\text{K}_3\text{Fe}(\text{CN})_6$  and 150 mM NaCl. Both GF anode and cathode were placed at 2.5 cm from the  
18 separator together with an Ag/AgCl/KCl sat. reference in each compartment (figure S1,  
19 Supplementary Information). Lactate was supplemented in the anodic compartment and  
20 interval between additions of 30 mM lactate was evaluated by current monitoring;  
21 decreasing of current corresponds to a decrease in lactate availability. To ensure sterility,  
22 all the reactors components were autoclaved at  $120 \text{ }^\circ\text{C}$  and then assembled in sterile  
23 conditions. The oxic condition was achieved during the whole experiment with vent-caps  
24 and sterile filters (pore size:  $0.2 \mu\text{m}$ ) to ensure atmospheric exchange.

25 To apply fixed potential at electrodes, the anode and the cathode were symmetrically and  
26 continuously polarized by applying well-defined potentials against a reference electrode  
27 (figure S1, Supplementary Information). The polarization effect was evaluated for anodic  
28 potential varying between  $-0.3 \text{ V}$  and  $+0.5 \text{ V}$  (vs. Ag/AgCl/KCl sat.) while the potential at  
29 the cathode was oppositely poised. Both the polarization step and the electrochemical  
30 characterizations were carried out in oxic condition.



1       HERE FIGURE 1

2  
3       In a second set of experiments, *S. oneidensis* bacteria harvested in the log phase and  
4 resuspended in MR1 medium containing 30 mM lactate were used and the anodic and  
5 cathodic applied voltage at the carbon felt electrodes were varied from - 0.3 V to + 0.5 V  
6 vs. Ag/AgCl/KCl sat. ( $E_c = - E_a$ ).

7       Chronoamperometry experiments were performed over 19 days (Figure 2). In the  
8 presence of a negative polarization, the current was 60  $\mu$ A after 1 day, dropped down to  
9 less than 10  $\mu$ A after 4 days and then continued to decrease despite the regular lactate  
10 feeding. In the absence of polarization, the initial current was smaller (*ca.* 35  $\mu$ A) and also  
11 underwent a rapid decrease but then increased again until it reached *ca.* 20  $\mu$ A after 19  
12 days. When a +0.3 V vs. Ag/AgCl/KCl sat. polarization was applied, the initial decrease  
13 phase was also observed and the current values varied around 50  $\mu$ A average value. Finally,  
14 for a + 0.5 V vs. Ag/AgCl/KCl sat. potential, the current was stable over the first days of  
15 the experiment and then increases progressively to reach *ca.* 120  $\mu$ A. Nevertheless, we  
16 observed a reproducible dispersion of current values at a given time points, suggesting that  
17 such high potentials may impact the bacterial activity and/or the stability of the bacteria-  
18 electrode interface.

19  
20       HERE FIGURE 2

21  
22       Cyclovoltamograms were recorded at different times of the experiments (Figure 3).  
23 After one day, regardless of the applied potential, cyclovoltamograms exhibited a pair of  
24 reversible faradaic peaks centered at - 0.4 V vs. Ag/AgCl/KCl sat. corresponding to  
25 riboflavin. An endogenous oxidation wave of similar shape and intensity was observed in  
26 the high positive and negative potential regions (Figure 3a) responsible for an extracellular  
27 electron transfer. After 4 days, the oxidation wave has increased in intensity for a + 0.5 V  
28 vs. Ag/AgCl/KCl sat. and a + 0.3 V vs. Ag/AgCl/KCl sat. polarization, in agreement with  
29 the chronoamperometry measurements (Figure 3b). In parallel, in the negative potential  
30 range, the signal becomes more complex, with an apparent splitting of the riboflavin peak.  
31 Such a splitting is confirmed at day 8 for all polarization conditions (Figure 3c). In parallel,



1 the oxidative wave has gained in intensity for systems under a positive polarization and  
2 especially at + 0.5 V vs. Ag/AgCl/KCl sat.. These two observations are still valid at day 19  
3 although, for a + 0.5 V vs. Ag/AgCl/KCl sat. polarization, the two peaks appear to have  
4 merged (Figure 3d), resulting in a single signal in the same potential region but much  
5 broader than at day 1 (Figure 3a).

6  
7 HERE FIGURE 3

8  
9 These sets of experiments confirm the existence of different electron transfer mechanism  
10 (DET and/or MET/riboflavin), depending on the applied potential. On the one hand, the  
11 continuous increase in the intensity of the signal in the positive potential range suggests  
12 that the DET mechanism becomes more and more effective with time, especially under  
13 highly positive polarization conditions. This should reflect that an increasing number of  
14 bacteria are in direct contact with the graphite felt electrodes (*i.e.* anode colonization)  
15 and/or that the electron transfer at the bacteria/electrode interface becomes facilitated. For a  
16 non-polarized electrode or a polarization at -0.3 V vs. Ag/AgCl/KCl sat. the increasing of  
17 the wave intensity is less marked. On the other hand, the signals in the negative potential  
18 range do not evolve much in intensity but rather in shape. This can indicate the production  
19 of other mediators by the bacteria and/or a modification of the mediator/graphite interface.  
20 In particular, the splitting of the riboflavin peak may correspond to the co-existence of  
21 molecules originating from bacteria in solution and within the deposited biofilm.

22 To clarify these points, SEM imaging of the samples were performed at the end of the  
23 chronoamperometry experiments (Figure 4). The lowest density of bacteria is observed at a  
24 negative potential ( $E_a = -0.3$  V vs. Ag/AgCl/KCl sat.). These bacteria are embedded in a  
25 dense biofilm. Under non-polarized condition or positive applied potential, a similar  
26 situation is observed but the cell density seems higher, although biological analyses would  
27 be necessary to quantitatively ascertain this point. While the specific case of negative  
28 polarization fits well with the measurement of a very low current value, it is clear that the  
29 biofilm morphology, as it is observed here, cannot explain the differences in the  
30 electrochemical behavior for the other polarization conditions. Hence, the applied potential  
31 should have a deep impact on the internal structure and organization of the biofilm,

1 influencing the diffusion of the mediator and/or the connectivity of the conductive  
2 elements.

3  
4 HERE FIGURE 4

5  
6 **Double compartment studies.** In a step forward the development of a complete MFC, a  
7 dual compartment reactor was set up using an ultrafiltration membrane (pore size = 0.2  $\mu\text{m}$ )  
8 as a separator to prevent bacterial diffusion from the anodic side to the cathodic  
9 compartment. A potential of either + 0.3 V or - 0.3 V was applied at the GF anode with an  
10 oppositely polarized cathode. The electrochemical behavior under symmetrical polarization  
11 was monitored by chronoamperometry and the evolution of the anode potential,  
12 polarization and the power curves as function of time were established

13 Figure 5 displays the evolution of the polarization and power curves over 19 days using a  
14 symmetrical polarization when the anode was polarized at + 0.3 V vs. Ag/AgCl/KCl sat.  
15 (vs.  $E_c = - 0.3$  V vs. Ag/AgCl/KCl sat.). At day 2, the potential of the bioanode is equal to -  
16 0.4 V vs. Ag/AgCl/KCl sat. and remains stable until the end of the experiment (Figure 5a).  
17 The anodic current density at  $E_a = 0$  V vs. Ag/AgCl/KCl sat. evolves from 70 (day 2) to 30  
18  $\text{mA}\cdot\text{m}^{-2}$  (day 19). The evolution of the MFC polarization curves (Figure 5b) is a result of  
19 the anodic and cathodic  $I$ - $V$  profiles. MFC OCV remains stable at *ca.* 0.65 V, indicating  
20 that the electroactivity established at the bioanode is stable. The internal resistance  
21 (polarization curve slope,  $R_{\text{INT}} = \Delta E/j$ ) increases with time. Since no detrimental clogging  
22 of the separator was observed after 19 days, such an increase indicates a modification of  
23 electron transfer at the surface of the bioanode. This is confirmed by evolution of the  
24 current density (the maximum current density decreases from *ca.* 100 to 35  $\text{mA}\cdot\text{m}^{-2}$ ) and  
25 the power curves (Figure 5c) where a loss in power from *ca.* 20 (day 2) to *ca.* 7  $\text{mW}\cdot\text{m}^{-2}$   
26 (day 19) was observed.

27  
28 HERE FIGURE 5

29  
30 The same experiment was performed with a - 0.3 V vs. Ag/AgCl/KCl sat. negative  
31 potential applied to the bioanode. Figure 6 summarizes the evolution of the characteristic

1 polarization and power curves. At the open-circuit, the potential of the bioanode (figure 6a)  
2 regularly decreases from - 0.25 V (day 2) to - 0.6 V vs. Ag/AgCl/KCl sat. (day 15). The  
3 MFC OCV shown in figure 6b increases from 0.5 V (day 2) to 0.85 V (day 15).  
4 Additionally, the maximal power density increases from 12 to 20 mW.m<sup>-2</sup> (figure 6c). The  
5 initial maximal density of current is about 100 mA.m<sup>-2</sup> and decreases to 60 mA.m<sup>-2</sup> at day 5  
6 and then increases again to *ca.* 70 mA.m<sup>-2</sup> (17% recovered) and remains stable until day 15.  
7 Noticeably, a loss of electrochemical performances was observed after 19 days that can be  
8 explained by the depletion in lactate due to the absence of feeding after day 13.

9  
10 HERE FIGURE 6

11  
12 These results indicate two different behaviors as function of the polarization condition  
13 applied. Figure 7 summarizes the different situations: for both negative (- 0.3 V) and  
14 positive (+ 0.3 V) polarization, the maximal MFC current decreases in the first 5 days from  
15 100 to about 60 mA.m<sup>-2</sup>. Then, the value remains stable or slowly decreases. For an applied  
16 potential of + 0.3 V vs. Ag/AgCl/KCl sat., the MFC OCV remains constant at + 0.65 V  
17 while the power density decreases with time from *ca.* 20 mW.m<sup>-2</sup> to 5 mW.m<sup>-2</sup>. This can be  
18 linked to a modification of the electron transfer from bacteria to graphite electrodes,  
19 correlated with the increase of the ohmic losses of the MFC. On the contrary, for applied  
20 potential of - 0.3 V vs. Ag/AgCl/KCl sat., an increase of the MFC OCV, from 0.55 V to  
21 0.85 V, and power density, from 12 mW.m<sup>-2</sup> to 20 mW.m<sup>-2</sup>, are observed, keeping day 19  
22 data aside. The progressive increase of MFC OCV suggests a longer phase of stabilization  
23 for the bacteria in contact with the graphite anode compared to the positive applied  
24 potential. Additionally, the increase in power density can be linked to a better transfer of  
25 electron from the bacteria to the GF fibers.

26  
27 HERE FIGURE 7

28  
29 The microstructure of the electrode after 19 days was studied by scanning electron  
30 microscopy (figure 8). For a potential applied at the bioanode of + 0.3 V vs. Ag/AgCl/KCl

1 sat., the bacteria can be distinctly observed at the surface of fibers and form a uniform  
2 layer. This bacterial layer is embedded in a thin and dense biofilm which forms a sheath  
3 wrapped around the fibers. In contrast, for potential applied at the bioanode of - 0.3 V vs.  
4 Ag/AgCl/KCl sat., a large quantity of biomass (EPS, bacteria, ...) occupies the porosity  
5 formed by the graphite fiber network. The fibers themselves are embedded in a thick and  
6 porous biofilm structure composed of a high density of bacteria and EPS.

7  
8 HERE FIGURE 8  
9

10 These observations can be related to the results of the single compartment experiments.  
11 Under positive polarization, DET becomes rapidly prevalent over MET, which can be  
12 correlated with an efficient colonization of the fiber surface by bacteria. In these conditions,  
13 as the biofilm thickens, the electron transfer may become less favorable, leading to a  
14 decrease in the electrochemical performance as observed for MFC power density. Yet, the  
15 MFC OCV remains stable meaning that the efficiency of the redox reaction and electron  
16 transfer at the fiber surface remains the same. In contrast, under negative polarization, both  
17 DET and MET are involved in the electron transfer, so that bacteria growth is favored both  
18 in solution, *i.e.* in the felt porosity, and on the fiber surface. In these conditions, the increase  
19 in the power density with time can be related to the simultaneous development of both  
20 populations. Moreover, the ability of the biofilm to bridge graphite fibers may provide  
21 additional conduction pathways. Yet it is interesting to point out that such bridging  
22 structures were not observed in single compartment experiments. This difference can be  
23 explained considering that, in the latter situation, the cathode is also in contact with the  
24 bacterial suspension and can interfere with its behavior. As a matter of fact, in these  
25 conditions, colonization of the graphite felt used as the counter-electrode could be observed  
26 by SEM (figure S3, Supplementary Information). On the contrary, in the double  
27 compartment configuration, the oppositely polarized electrode is isolated from the bacterial  
28 anolyte by the ultrafiltration membrane so that bacterial behavior must be influenced only  
29 by the anodic potential of polarization.

## 30 31 CONCLUSION

1 In this work, the impact of polarization on the colonization of graphite felts used as  
2 bioanodes by wild type *S. oneidensis* in oxic conditions and its influence on the  
3 electrochemical performances of single- and double-compartment fuel cells were studied.  
4 In single-compartment MFC configurations, both anodic current and electrode colonization  
5 are directly correlated to positive polarization (+ 0.3 and + 0.5 V vs. Ag/AgCl/KCl sat.).  
6 Under non-polarized condition, a lower and less reproducible current is measured but it  
7 remains higher than the current measured under negative polarization. These results suggest  
8 that positive polarizations favor the felt colonization but also impact the properties of the  
9 resulting biofilm. However, we also observed an unintended colonization of the counter  
10 electrode, which suggests a combined effect of both fixed anode and cathode potentials. To  
11 elucidate this phenomenon and prevent the cathode colonization by bacteria, dual-  
12 compartments MFC experiments were carried out in the same polarization conditions. Two  
13 distinct behaviors are observed: (i) under positive polarization, MFC performance  
14 (polarization and power curves), anodic OCP and MFC OCV are already at their maximal  
15 value after a short time of polarization; however, MFC performances decrease rapidly with  
16 time. The biofilm appears thin and dense around the felt fibers. (ii) Under negative  
17 polarization, the electrochemical parameters progressively increase to reach their maximal  
18 values after more than 20 days. In this condition, the biofilm is thick, porous and fills the  
19 porosity of the felt. These two behaviors seem in agreement with the DET/MET hypothesis.  
20 At + 0.3 V vs. Ag/AgCl/KCl sat., the electrode colonization is controlled by DET leading  
21 to the accumulation of bacteria on the fiber surface. On the contrary, at - 0.3 V vs.  
22 Ag/AgCl/KCl sat., the MET mechanism is more effective, driving the development of  
23 bacteria in the liquid phase, *i.e.* colonization of the felt porosity. Yet, biofilm formation of  
24 the fiber surface is also observed, suggesting that DET can also occur. Whether a negative  
25 potential favor MET, hinder DET or both remains an open question.

26  
27

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14    **ABBREVIATIONS**

15    GF: graphite felt, Ea: anodic potential, Ec: cathodic potential, OD<sub>600</sub>: Optical Density at

16    600 nm, MFC: microbial fuel cell, OCV: open-circuit voltage, OCP: open-circuit potential,

17    SEM: scanning electron microscopy, cfu.mL<sup>-1</sup>: colony-forming unit per milliliter, I-V

18    curve: current-tension curve.

19

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## FIGURES CAPTIONS

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23 **Figure 1.** Cyclic voltammetry at  $10 \text{ mV}\cdot\text{s}^{-1}$  of a highly concentrated suspension of *S.*  
24 *oneidensis* in a PBS buffer supplemented with 30 mM lactate in oxic conditions. The  
25 potential range corresponding to mediated electron transfer (MET, red zone) and direct  
26 electron transfer (DET, blue zone) and the corresponding scheme are adapted from Roy *et*  
27 *al.* <sup>9</sup>.

1 **Figure 2.** Chronoamperometry measurements for *S. oneidensis*/graphite felt bioanodes in  
2 single-compartment reactors with an applied potential of (orange circles and squares) - 0.3  
3 V, (blue circles and squares) 0 V, (green circles and squares) + 0.3 V and (dark circles and  
4 squares) + 0.5 V

5 **Figure 3.** Cyclovoltammograms at  $10 \text{ mV}\cdot\text{s}^{-1}$  for *S. oneidensis*/graphite felt bioanodes in  
6 single-compartment reactors with an applied potential of (orange line) - 0.3 V, (blue line)  
7 0 V, (green line) + 0.3 V and (dark lines) + 0.5 V after (a) 1 day, (b) 4 days, (c) 8 days and  
8 (d) 19 days of polarization. Reference signal of GF/MR1 without bacteria and reference  
9 signal of riboflavin on figure S2 (Supplementary Information).

10 **Figure 4.** SEM observations of *S. oneidensis*/graphite felt bioanodes in single-  
11 compartment reactors. Representative morphology of colonized felts (a) and higher  
12 magnification images for bioanodes polarized at (b) + 0.5 V, (c) + 0.3 V, (d) non polarized,  
13 (e) - 0.3 V, after 19 days in working condition.

14 **Figure 5.** Electrochemical characterization of *S. oneidensis*/graphite felt//  
15  $\text{K}_3\text{Fe}(\text{CN})_6$ /graphite felt double-compartment reactor with a bioanode polarized at + 0.3 V  
16 after (orange circle) 2 days, (grey circle) 5 days, (green circle) 6 days, (blue circle) 15 days  
17 and (dark circle) 19 days: (a) anodic I-V, (b) polarization and (c) power curves.

18 **Figure 6.** Electrochemical characterization of *S. oneidensis*/graphite felt//  
19  $\text{K}_3\text{Fe}(\text{CN})_6$ /graphite felt double-compartment reactor with a bioanode polarized at - 0.3 V  
20 after (orange circle) 2 days, (grey circle) 5 days, (green circle) 6 days, (blue circle) 15 days  
21 and (dark circle) 19 days: (a) anodic I-V, (b) polarization and (c) power curves.

22 **Figure 7.** Temporal evolution of (dark circle) open-circuit voltage (OCV), (red circle)  
23 maximal power density and (blue circle) maximal current density for a *S.*  
24 *oneidensis*/graphite felt// $\text{K}_3\text{Fe}(\text{CN})_6$ /graphite felt double-compartment reactor with a  
25 bioanode polarized at (a) + 0.3 V and (b) -0.3 V.

26 **Figure 8.** SEM images of the anodic graphite felt interior after 19 days of function for a *S.*  
27 *oneidensis*/graphite felt// $\text{K}_3\text{Fe}(\text{CN})_6$ /graphite felt double-compartment reactor with a  
28 bioanode polarized at (a) + 0.3 V and (b) -0.3 V.

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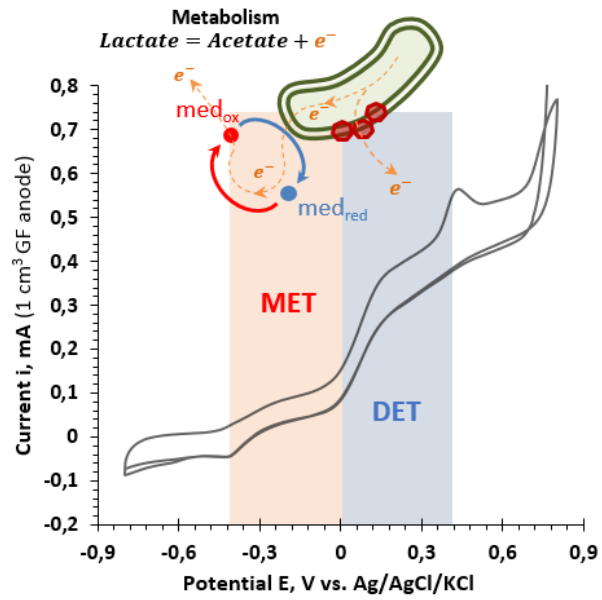
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# FIGURES

## FIGURE 1

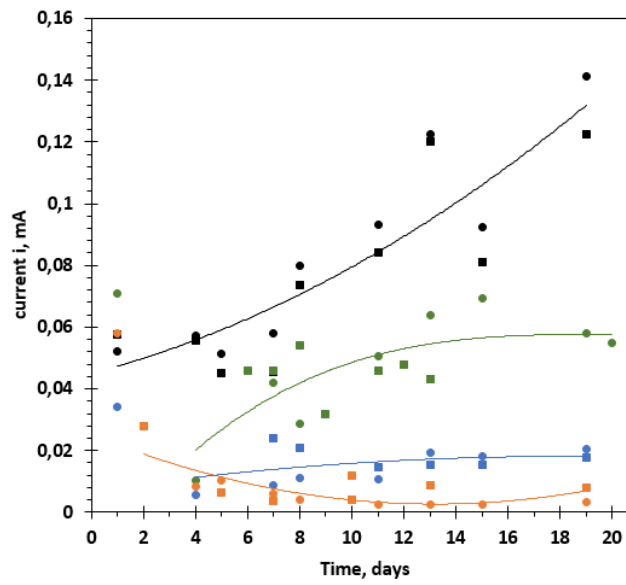


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## FIGURE 2



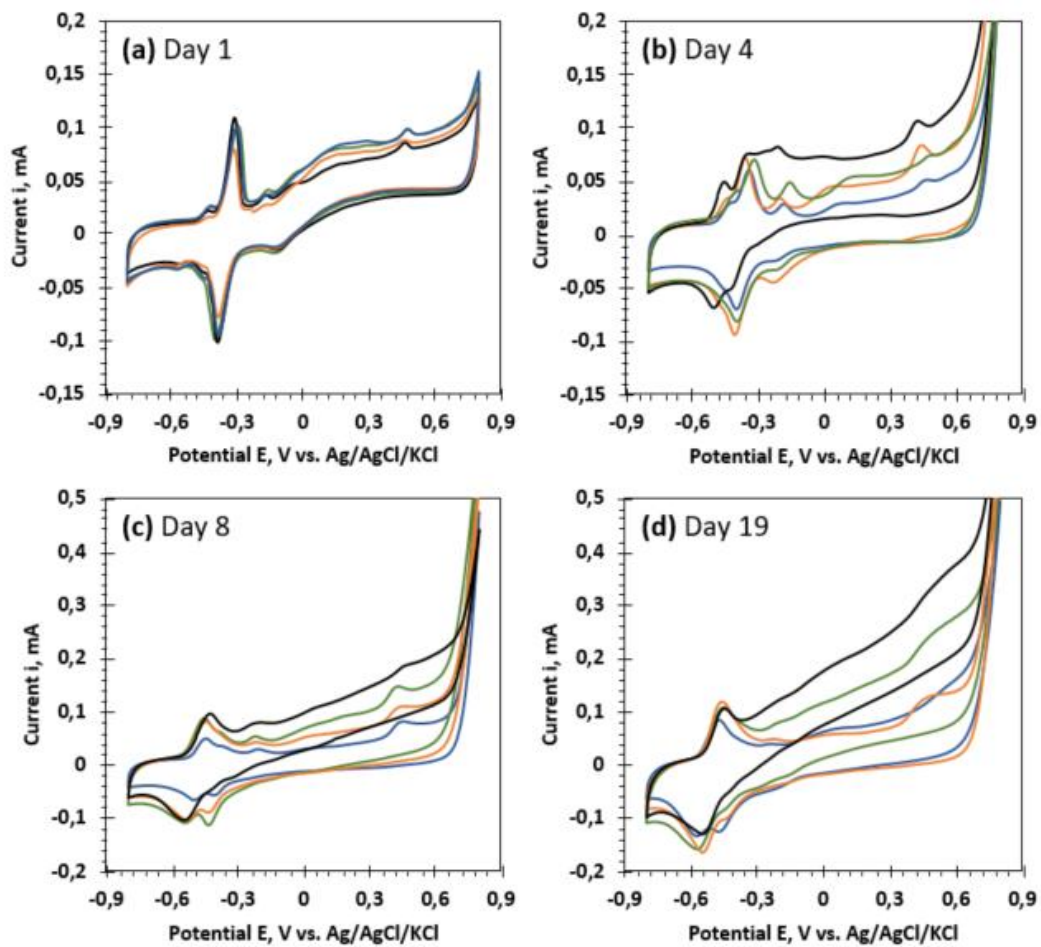
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FIGURE 3



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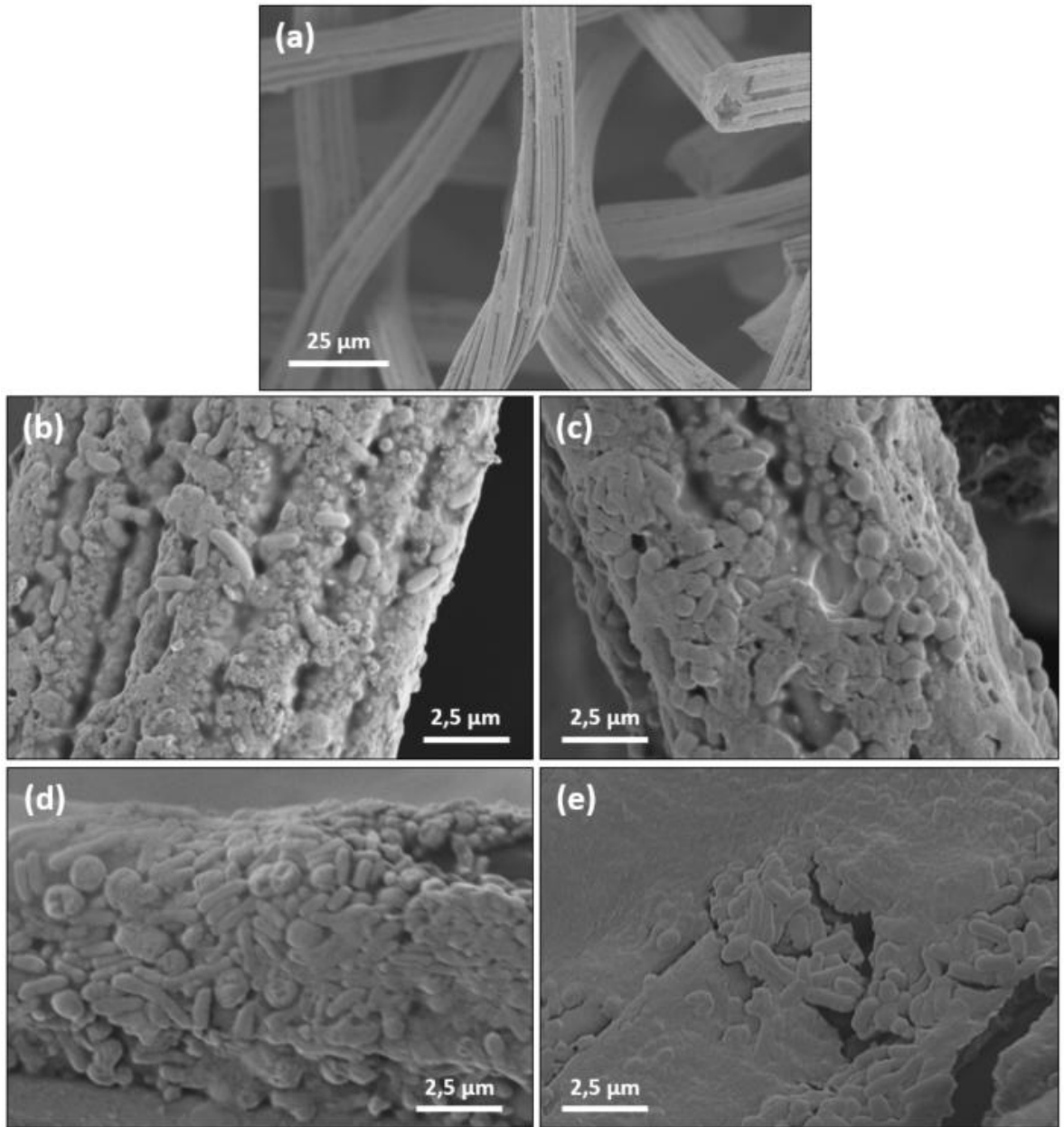
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FIGURE 4



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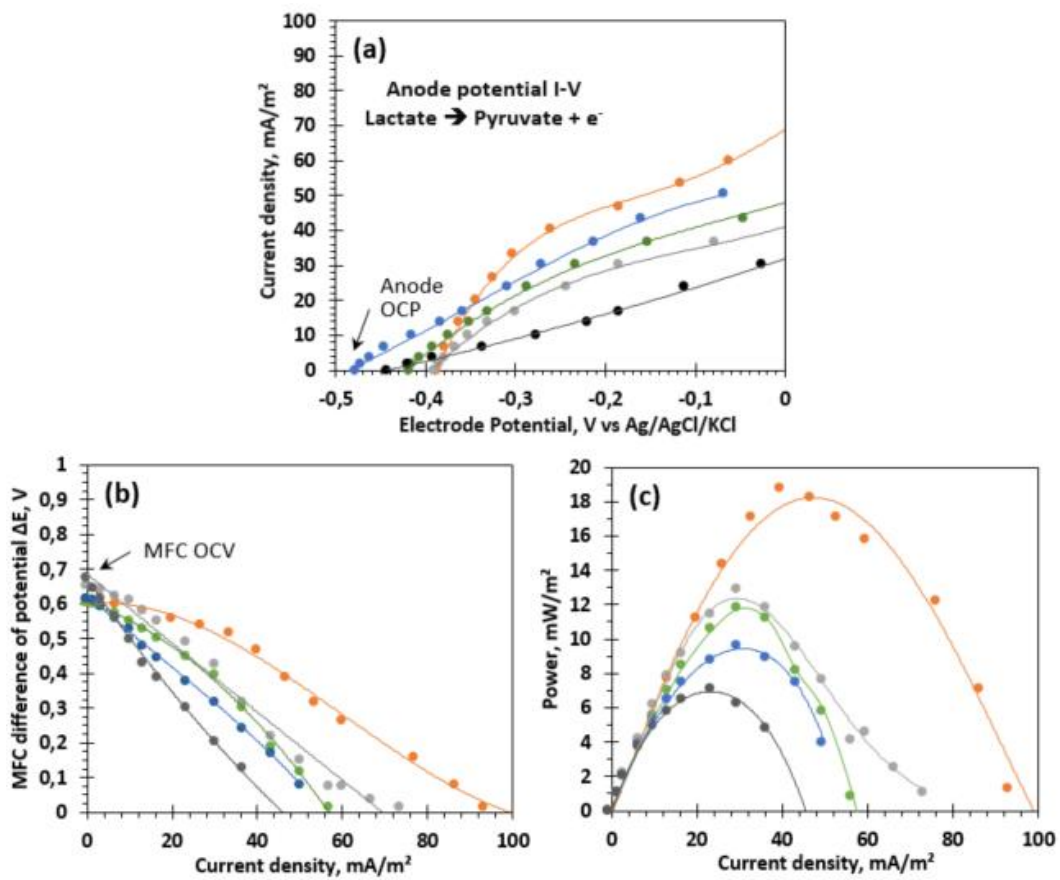
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FIGURE 5



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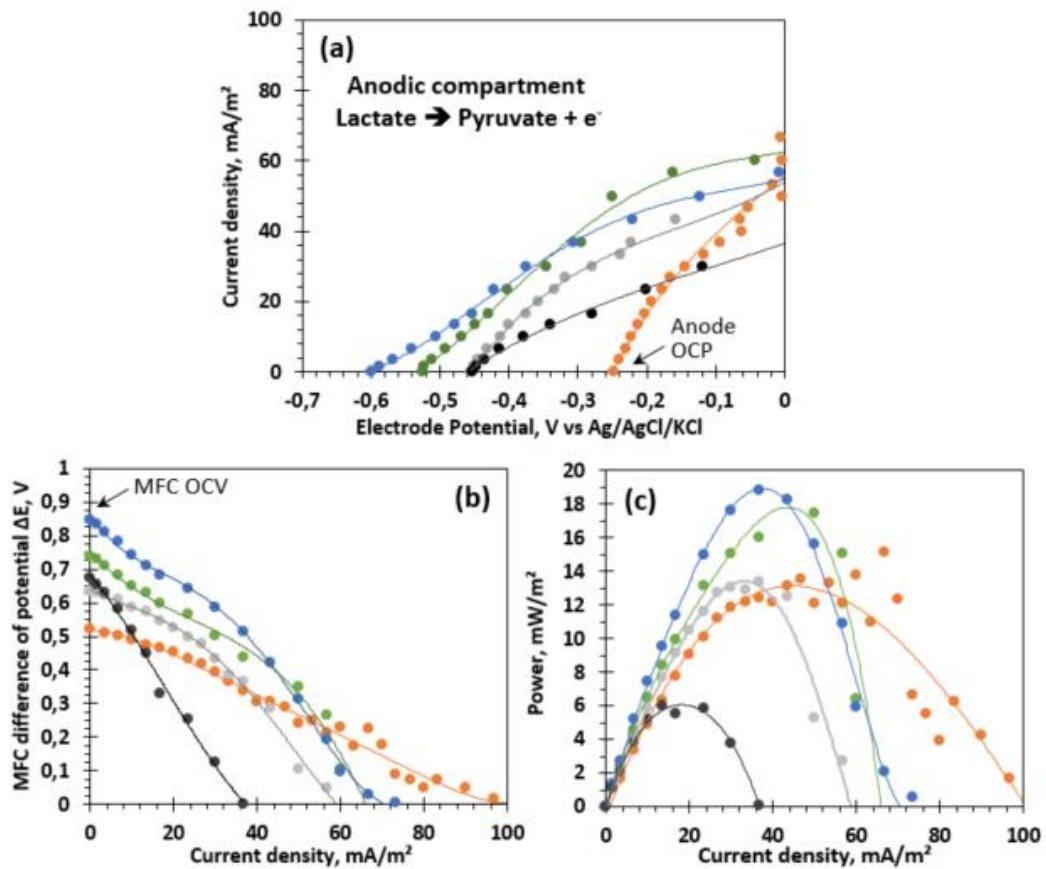
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FIGURE 6



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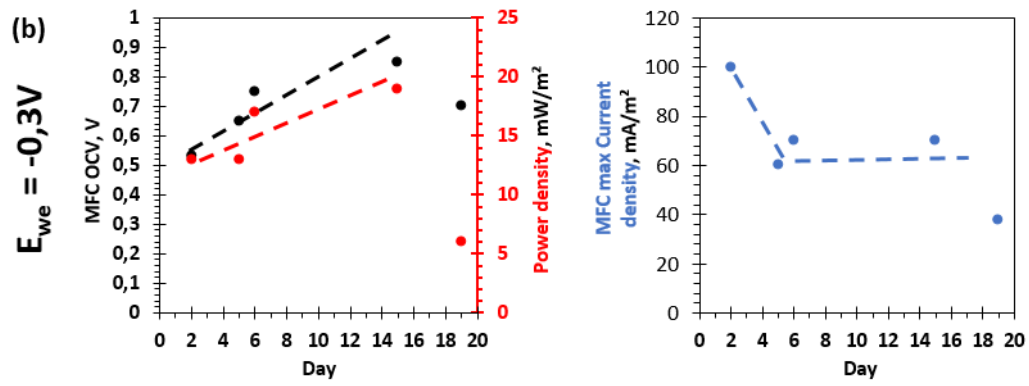
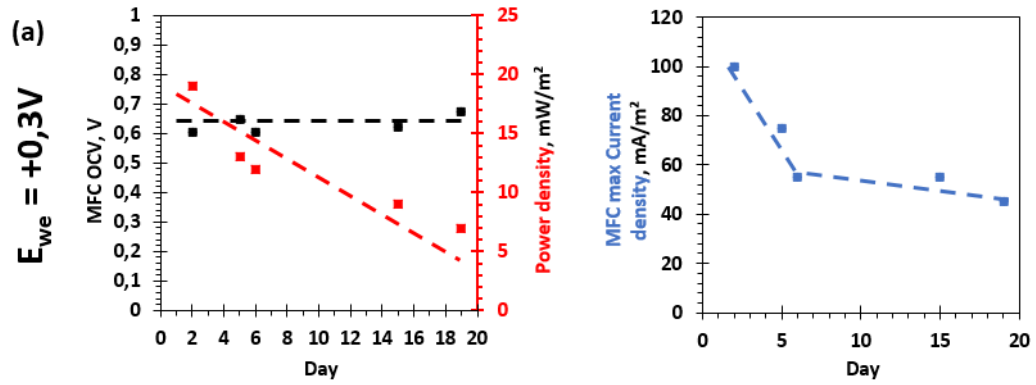
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FIGURE 7

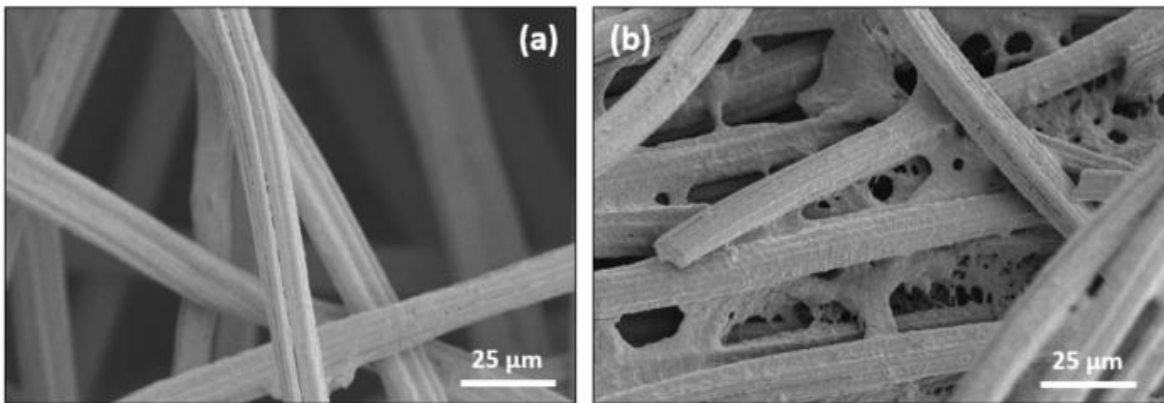


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FIGURE 8



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