## **Electronic supplementary information (ESI)**

#### Biofilm as a redox conductor: a systematic study of moisture and 1 2 temperature dependence of its electrical properties

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#### Sample preparations 16

17 Commercially available IDAs were used in this study (Fig. S1a; ALS IDA electrodes 18 model 012125) and consisted of 65 pairs of parallel gold rectangular bands, each 2 mm long x 10 µm wide x 90 nm thick, patterned onto quartz substrates. Electrode bands were separated 19 by a 5 µm gap and alternating electrodes were attached to large electrode contacts on opposite 20 sides of the array. 21

22 Geobacter sulfurreducens samples were prepared as reported previously.<sup>1,2</sup> Briefly, G. sulfurreducens biofilms were grown on electrodes poised at +0.5 V (vs. SHE) in freshwater 23 medium (ATCC 2260) with 10 mM acetate and excluding fumarate until the current output 24 stabilized. The water jacketed reactor (Pine Instruments, USA) was stirred and the 25 temperature was maintained at 30°C throughout growth using a temperature controlled water 26 bath. The system was maintained under anaerobic conditions by continuously sparging with a 27 80% N<sub>2</sub>/ 20% CO<sub>2</sub> gas mixture. Electrochemical gating measurements and cyclic 28 voltammetry were performed on the electrodes with a bipotentiostat (Pine Instruments, USA) 29 using the provided software to ensure that the system was performing similarly to samples 30 previously generated.<sup>1</sup> Data is shown in Figure S1. Before the electrode was removed from 31 the reactor, the gate potential,<sup>3,4</sup> defined as the average of the potentials applied to the two 32 electrodes while maintaining a fixed voltage offset of 0.1 V (source-drain voltage) between 33 the electrodes, was set to the potential where maximum conducted current was observed (-34 0.19 V vs. SHE) for 3 minutes. This was done with the intention of preparing biofilms with 35 equal concentrations of reduced and oxidized redox cofactors for maximum rate of electron 36 transport.<sup>3,4</sup> However, it is unclear if this state remained intact after removing the biofilm 37 from the electrolyte prior to analysis. Further characterization of the concentrations of reduced 38 and oxidized cofactors during electrochemical tests was beyond the scope of this study. The 39 electrode was then removed, rinsed 3 times in DI water and placed in an anaerobic chamber 40

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#### **Electronic supplementary information (ESI)**

41 (relative humidity ~15%) overnight to facilitate drying of the biofilm. The biofilm was dried

- 42 in order to preserve the state of the biofilm.
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45 Figure S1. Source-drain current vs. gate potential dependencies generated by electrochemical
46 gating experiments of biofilms used in this study under *in situ* conditions prior to being
47 removed from medium for subsequent *ex situ* measurements. The current magnitudes are
48 consistent to what has been previously observed for the same experimental parameters.<sup>3</sup> the
49 variation in current magnitude among biological replicates is not uncommon under in situ
50 conditions. However, the difference in conducted current in the reactor medium did not affect
51 results during hydration tests (Fig S5b).

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53 Shewanella oneidensis MR-1 biofilms were grown in microbial three-electrode electrochemical cells similar to those previously reported,<sup>5,6</sup> with the IDA serving as the 54 working electrode poised at 0.3 V vs. Ag/AgCl (0.5 V vs. SHE). In order to culture the 55 bacteria for these reactors, S. oneidensis MR-1 was struck out on LB agar plates from frozen 56 bacterial stock and incubated at 30 °C overnight to isolate single colonies. Biological replicate 57 cultures were grown by selecting morphologically similar colonies with a sterile loop to 58 inoculate anaerobic (100% N<sub>2</sub> atmosphere) modified M1 medium<sup>6-8</sup> containing 20 mM Na-59 (L)-lactate as donor and 20 mM Na-fumarate as acceptor. After 24 hours of incubation with 60 shaking in a temperature controlled chamber at 30 °C, a consistent OD<sub>600</sub> of ~0.16 was 61 reached (i.e.  $1.6 \times 10^8$  cfu/mL using the previously determined conversion factor ~1  $\times 10^9$ 62 cfu/mL/OD<sup>6</sup>). These stationary phase cultures (fumarate completely consumed) may then be 63 used to inoculate replicate devices as desired. 64

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Poly(N-vinylimidazole [Os(bipyridine)<sub>2</sub>Cl])<sup>+/2+</sup> (referred to here as PVI-Os(bipy)<sub>2</sub>Cl) 65 was formed on IDAs using established literature methods<sup>9</sup> in which a 1:0.22:0.44 solution of 66 PVI-Os(bipy)<sub>2</sub>Cl (10 mg/L):PEGDE crosslinker (3 mg/mL): ethanol (100%) was mixed 67 together and drop cast to cover the electrode surface. The electrode was then allowed to dry 68 overnight in a dessicator. Polyaniline (PANI) was electropolymerized on IDAs using 69 established literature procedures<sup>10</sup> in which 50 mM of aniline monomer in 0.5 M H<sub>2</sub>SO<sub>4</sub> was 70 electrodeposited by first sweeping the potential from open circuit potential (0.57 V) to 1 V vs. 71 SHE and then holding the potential until 2.5 C/cm<sup>2</sup> of charge had passed through the electrode 72 to obtain a film ~25  $\mu$ m thick. The electrode was then rinsed in clean 0.5 M H<sub>2</sub>SO<sub>4</sub> before use. 73 A BioLogic potentiostat Model VMP3 (BioLogic, Inc.) and platinum counter electrode were 74 75 used to electropolymerize PANI. At pH=0, the electropolymerized PANI films were expected 76 to be fully protonated (doped) and metallic, whose conductivity negligibly depends on humidity (as we observed in our experiments).<sup>11,12</sup> It should be noted that PANI film's 77 conducivity can change with humidity depending on the polymer synthesis and the film 78 preparation.<sup>13,14</sup> 79

#### 80 Two-point probe measurements and conductivity calculation

In a typical steady state current voltage measurement, biases from 0 to 2 V were swept between two contacts of the IDA (Fig. S1a) and currents were measured. The conductance of the films, G, was calculated from the slope of current vs. voltage (Fig. S1b). Conductivity was then calculated from the conductance and the geometrical factor S,<sup>1,15</sup> which was calculated in

85 our previous work to be  $20.4.^{1}$ 

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87 Figure S2. (a) IDA electrode (<u>http://www.als-japan.com/1379.html#defaultTab13</u>)
88 dimension; (b) a typical current-voltage (I-V) curve measured for a *G. sulfurreducens* biofilm
89 or Os-complex film showing the linear relationship between current and applied bias; (c, d)

90 Chemical structures of the osmium redox complex and doped polyaniline, respectively.

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92 Figure S3. Film conductivity as a function of water content at 20 C controlled by changing
93 the relative humidity in the measuring chamber. The conductivity measurements were
94 reproduced for two independent films and two replicates for each film.





96 **Figure S4**. (a) Current-voltage (I-V) curves of *G. sulfurreducens* biofilm (T=25C, RH=45%), 97 Osmium redox polymer film (T=25C, RH=35%) and polyaniline film (T=25C, RH=35%); (b) 98 I-V curves of *S. oneidensis* MR-1 biofilm measured at 25 C and relative humidity from 45% 99 to 85%; and bare IDA electrode at T=25C, RH=38.5%. The current level of bare IDA-100 electrode (noise current) does not discernably change at different temperature and humidity.

### **Electronic supplementary information (ESI)**

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(a) *G. sulfurreducens* Biofilm (25 °C, RH = 95.0%)

(b) 25 °C, RH = 95.0%





103 Figure S5. Two-point probe transient current measurements. (a) G. sulfurreducens biofilm at

- 104 T = 25 °C, RH = 95.0% at two different biases with opposite polarity. (b, c, d) G.
- 105 sulfurreducens biofilm, Os-complex film and S. oneidensis biofilm at different temperature

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106 and humidity conditions: (b) T = 25 °C, RH = 95.0%, (c) T = 25 °C, RH = 38.7% and (d) T = 107 12 °C, RH = 86.8%. The lines connecting the data points are plotted for visual aid.



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109 Figure S6. (a) Conductivity of Os-complex film in ambient air at constant RH = 60% and

110 temperatures from 15 °C to 35 °C. Water content was calculated from relative humidity and

- 111 temperature using this website: http://www.owlstonenanotech.com/humidity/calculator. (b)
- 112 Conductivity of Os-complex film in ambient air at fixed water content 12,000 ppm and
- 113 temperatures from 12 °C to 35 °C.
- 114

#### 115 **Reference**

- 116 1 M. D. Yates, J. Golden, J. Roy, S. M. Strycharz-Glaven, S. Tsoi, J. Erickson, M. Y. El-
- 117 Naggar, S. C. Barton and L. Tender, *Phys. Chem. Chem. Phys.*, 2015.
- 118 2M. D. Yates, J. P. Golden, S. M. Strycharz-Glaven, S. D. Tsoi, J. S. Erickson, M. Y. El-
- 119 Naggar, S. A. Calabrese Barton and L. M. Tender, *Nat Nano*, 2016, In review.
- 120 3 M. D. Yates, J. P. Golden, J. Roy, S. M. Strycharz-Glaven, S. Tsoi, J. S. Erickson, M. Y. El-
- 121 Naggar, S. C. Barton and L. M. Tender, *Phys. Chem. Chem. Phys.*, 2015, 17, 32564–32570.
- 4R. M. Snider, S. M. Strycharz-Glaven, S. D. Tsoi, J. S. Erickson and L. M. Tender, *Proc. Natl. Acad. Sci.*, 2012, **109**, 15467–15472.
- 5E. Marsili, J. B. Rollefson, D. B. Baron, R. M. Hozalski and D. R. Bond, *Appl. Environ. Microbiol.*, 2008, 74, 7329–37.
- 126 6N. D. Kirchhofer, X. Chen, E. Marsili, J. J. Sumner, F. W. Dahlquist and G. C. Bazan, *Phys.* 127 *Chem. Chem. Phys. PCCP*, 2014, **16**, 20436–43.
- 7B. Cao, L. Shi, R. N. Brown, Y. Xiong, J. K. Fredrickson, M. F. Romine, M. J. Marshall, M.
  S. Lipton and H. Beyenal, *Environ. Microbiol.*, 2011, 13, 1018–31.
- 130 8B. Cao, P. D. Majors, B. Ahmed, R. S. Renslow, C. P. Silvia, L. Shi, S. Kjelleberg, J. K.
- 131 Fredrickson and H. Beyenal, *Environ. Microbiol.*, 2012, **14**, 2901–10.

## **Electronic supplementary information (ESI)**

- 132 9J. W. Gallaway and S. A. Calabrese Barton, J. Am. Chem. Soc., 2008, 130, 8527-8536.
- 133 10 E. W. Paul, A. J. Ricco and M. S. Wrighton, J. Phys. Chem., 1985, 89, 1441–1447.
- 134 11 A. G. Macdiarmid, J. C. Chiang, A. F. Richter and A. J. Epstein, *Synth. Met.*, 1987, **18**, 135 285–290.
- 136 12 A. J. Epstein, J. M. Ginder, F. Zuo, R. W. Bigelow, H.-S. Woo, D. B. Tanner, A. F.
- 137 Richter, W.-S. Huang and A. G. MacDiarmid, Synth. Met., 1987, 18, 303–309.
- 138 13 J. P. Travers and M. Nechtschein, *Synth. Met.*, 1987, **21**, 135–141.
- 139 14 T. Taka, Synth. Met., 1993, **57**, 5014–5019.
- 140 15 J. Kankare and E.-L. Kupila, J. Electroanal. Chem., 1992, **322**, 167–181.
- 141

142