Syntrophic anaerobic photosynthesis via direct interspecies electron transfer

Phuc T. Ha¹, Stephen R. Lindemann², Liang Shi³, Alice C. Dohnalkova⁴, James K. Fredrickson², Michael T. Madigan⁵ & Haluk Beyenal¹

Microbial phototrophs, key primary producers on Earth, use H₂O, H₂, H₂S and other reduced inorganic compounds as electron donors. Here we describe a form of metabolism linking anoxygenic photosynthesis to anaerobic respiration that we call ‘syntrophic anaerobic photosynthesis’. We show that photoautotrophy in the green sulfur bacterium Prosthecochloris aestuarii can be driven by either electrons from a solid electrode or acetate oxidation via direct interspecies electron transfer from a heterotrophic partner bacterium, Geobacter sulfurreducens. Photosynthetic growth of P. aestuarii using reductant provided by either an electrode or syntrophy is robust and light-dependent. In contrast, P. aestuarii does not grow in co-culture with a G. sulfurreducens mutant lacking a trans-outer membrane porin-cytochrome protein complex required for direct intercellular electron transfer. Syntrophic anaerobic photosynthesis is therefore a carbon cycling process that could take place in anoxic environments. This process could be exploited for biotechnological applications, such as waste treatment and bioenergy production, using engineered phototrophic microbial communities.

¹The Gene and Linda Voiland School of Chemical Engineering and Bioengineering, Washington State University, Pullman, Washington 99164, USA.
²Biological Sciences Division, Pacific Northwest National Laboratory, Richland, Washington 99354, USA.
³Department of Biological Sciences and Technology, School of Environmental Studies, China University of Geoscience, Wuhan, Hubei 430074, China.
⁴Environmental Molecular Sciences Laboratory, Pacific Northwest National Laboratory, Richland, Washington 99354, USA.
⁵Department of Microbiology, Southern Illinois University, Carbondale, Illinois 62901, USA.

Correspondence and requests for materials should be addressed to H.B. (email: beyenal@wsu.edu).
Photosynthetic CO₂ fixation by cyanobacteria (oxygenic phototrophs), and purple and green sulfur bacteria (anoxygenic phototrophs) accounts for nearly one-half of global primary productivity. Major electron donors for anoxygenic photosynthesis—an anaerobic process—include reduced inorganic compounds such as H₂S, S⁰, H₂, and Fe²⁺ (2). In anoxic environments, electrons flow between microbes that form intimate syntrophic consortia to support complementary metabolisms. Perhaps the most intriguing of these electron flow mechanisms is the discovery of direct interspecies electron transfer, a process in which electrons are carried between cells through physical contact of conductive electron carriers. Some examples include anaerobic ethanol oxidation by consortia composed of two Geobacter species and the conversion of ethanol to methane by co-cultures of Geobacter species, and methanogenic Archaea such as Methanoseta. Although studies have suggested that direct interspecies electron transfer is common in anoxic methanogenic environments, whether related phenomena are more widespread, including in illuminated anoxic environments, is unknown. Here, we investigate this possibility and report the discovery of a new form of syntrophic metabolism, syntrophic anaerobic photosynthesis. We show that photoautotrophy in the green sulfur bacterium Prosthecochloris aestuarii can be driven by either electrons from a solid electrode or acetate oxidation via direct interspecies electron transfer from a heterotrophic partner bacterium, Geobacter sulfurreducens. Both interspecies electron transfer between G. sulfurreducens and P. aestuarii, and electron uptake by P. aestuarii are light-dependent processes. Light is required to drive photosynthetic metabolism in the phototroph. The discovery of syntrophic anaerobic photosynthesis reveals new possibilities for bioengineering phototrophic microbial communities for applications in the areas of waste treatment and bioenergy production.

Results

Electrode as electron donor for P. aestuarii. Until now, the purple non-sulfur bacterium Rhodopseudomonas palustris TIE-1 has been the only phototroph known to be able to assimilate electrons from a solid electrode. A few studies also suggested that some anaerobic phototrophs in a mixed culture are capable of using an electrode as an electron donor for photosynthesis. However, neither the mechanisms nor the phototrophs responsible for electron exchange with electrodes have yet been identified, leaving an open question of whether other phototrophs can also capture electrons directly from a solid electrode for photosynthesis. In a previous study on electron transfer in a microbial mat harvested from Hot Lake—an episomitic lake in northern Washington (WA, USA)—we found the green sulfur bacterium Prosthecochloris aestuarii to predominate at the tip of a carbon microelectrode inserted into the illuminated mat. In subsequent experiments, a pure culture of P. aestuarii was isolated. P. aestuarii can grow photoautotrophically utilizing sulfide or elemental sulfur (S⁰) as an electron donor. Recent studies have described several sulfur-oxidizing bacteria that are also capable of donating/accepting electrons from an electrode. To determine whether the Hot Lake P. aestuarii can also utilize electrons from a solid electrode, we inoculated it into a sterile, two-chamber bioelectrochemical system containing growth medium in which CO₂ was the sole carbon source with it. The electrode was polarized at −600 mVAg/AgCl (slightly lower than the redox potential of H₂S (E⁰[H₂S/H2S] − 479 mVAg/AgCl)), and electron transfer from the electrode was assessed as cathodic current. A light-dependent cathodic current was detected that gradually increased over 24 h; after 17 days, the current reached 104 μA (~36.74 μA cm⁻²) (Fig. 1a). This electron uptake rate with a phototroph is well above that previously achieved with R. palustris TIE-1 (1.5 μA cm⁻²). The continued development of cathodic current after medium exchanges indicates that no soluble electron shuttles are required for electron uptake by P. aestuarii (Supplementary Fig. 1a).

Scanning electron microscopy of a section of the electrode confirmed that the material accumulating was indeed a biofilm of P. aestuarii cells (Fig. 1b–d) in intimate contact with the electrode surface. These electrode-attached cells were used to inoculate fresh P. aestuarii growth medium containing sulfide as an electron donor, and growth occurred, confirming their viability on the electrode (Supplementary Fig. 1b).

To examine whether molecular hydrogen (H₂) mediated electron transfer between the electrode and P. aestuarii, we inoculated medium containing H₂ as the sole electron donor under illumination with P. aestuarii. P. aestuarii did not grow under these conditions, even after prolonged incubation (>10 days), indicating that this phototroph cannot use H₂ as an electron donor for photosynthesis (Supplementary Fig. 2a) and that uptake of electrons from the electrode was not mediated by H₂.

Co-culture of P. aestuarii with G. sulfurreducens. Bolstered by our findings with electrode-dependent photosynthesis by P. aestuarii (Fig. 1), we investigated the possibility that this organism could be grown in co-culture with a heterotrophic partner bacterium. In addition to transferring electrons to other bacterial species, the anaerobic and chemoheterotrophic bacterium Geobacter sulfurreducens can donate electrons to a solid electrode. Since our experiments showed that P. aestuarii grew photoautotrophically using the electrons supplied by an electrode (Fig. 1), we hypothesized that G. sulfurreducens could also donate electrons to cells of
P. aestuarii to support the growth of both species under conditions in which neither could grow independently. To test this, we cultivated the two organisms in an illuminated, anoxic medium containing acetate but devoid of HS\(^-\), S\(^0\) or any other potential electron donors or acceptors. A small amount of thiosulfate (0.5 mM) was added as a source of sulfur for biosynthetic needs; however, thiosulfate is used neither as a photosynthetic electron donor by P. aestuarii\(^1,2,20\) nor as an electron acceptor by G. sulfurreducens\(^21\). Since P. aestuarii is unable to fix CO\(_2\) in the absence of HS\(^-\) or S\(^0\) (ref. 14) and G. sulfurreducens is unable to oxidize acetate in the absence of an electron acceptor, we reasoned that if a co-culture containing both organisms grew, it must be the result of interspecies electron transfer.

In these co-culture experiments, growth did indeed occur concomitantly with acetate consumption (Fig. 2a,b). Both P. aestuarii and G. sulfurreducens grew within the co-culture, as evidenced both by flow cytometry (Supplementary Fig. 3) and electron microscopy (Fig. 2c–e), in which the morphologically distinct species could be quantified. The TEM whole mount images of P. aestuarii (P.a) and G. sulfurreducens (G.s) co-cultures also displayed intimate cell connections, possibly functioning in electron transfers from G. sulfurreducens to P. aestuarii (Fig. 2d,e). The changes of acetate consumption and cell density when switching the co-cultures from dark to illuminated condition and vice versa (Fig. 3) suggest that the syntrophic growth between G. sulfurreducens and P. aestuarii is strictly light-dependent.

When these co-cultures were subcultured in fresh medium containing acetate, they continued to grow and consume acetate (Supplementary Fig. 4). This indicates that the syntrophic interaction was stable over multiple generations. The slower increase of biomass in subcultures than in original co-cultures is likely due to the cell density in the co-culture inoculum being lower than that in the two pure cultures. We conclude that growth of the co-culture represents a previously unrecognized form of anoxygenic photosynthesis that exploits the complementary metabolisms of a heterotroph and a phototroph to the benefit of both. Because growth of the co-culture strictly depends on the metabolic activities of each partner bacterium, we term this process as 'syntrophic anaerobic photosynthesis'.

Direct electron transfer supports syntrophic anaerobic photosynthesis. Although our co-culture experiments revealed a metabolic interdependency between a chemotroph and a photo-troph, the mechanism of electron transfer remained unclear. Chemically mediated interspecies electron transfer via exchange of a soluble organic compound such as formate, or of H\(_2\), was a possible mechanism for the transfer of electrons from G. sulfurreducens to P. aestuarii. However, this was eliminated as a possibility in our co-cultures because P. aestuarii is unable to use H\(_2\) or formate as an electron donor (Supplementary Fig. 2a). Three additional lines of evidence support our conclusion that the metabolic interdependency between G. sulfurreducens and P. aestuarii is not the result of a soluble electron carrier but instead requires cell–cell contact and direct interspecies electron transfer. First, the two organisms did not grow when cultured in the same medium but physically separated into two chambers by membrane filters (0.1-μm pore size), whereas a parallel culture containing both organisms in the same reactor grew (Supplementary Fig. 5). Second, in contrast to co-cultures with wild-type G. sulfurreducens under identical conditions, co-cultures of P. aestuarii and a G. sulfurreducens deletion mutant lacking the ombB-omaB-omcB-orfS-ombC-omaC-omcC gene cluster did not grow and acetate was not consumed (Fig. 4). The ombB-omaB-omcB-orfS-ombC-omaC-omcC gene cluster encodes a trans-outter membrane porin-cytochrome protein complex essential for extracellular electron transfer by G. sulfurreducens to solid electron acceptors such as ferrihydrite\(^22\) or a solid electrode (Fig. 4a). Kanamycin (200 μg ml\(^{-1}\)) was added to the culture in...

**Figure 2 | Syntrophic growth of G. sulfurreducens and P. aestuarii.** Variation in (a) cell density and (b) acetate consumption over time of P. aestuarii and G. sulfurreducens in axenic cultures and co-cultures. Each symbol is mean ± s.d. (n = 3). (c) SEM image of co-culture (scale bar, 5 μm). (d,e) TEM whole mount images of G. sulfurreducens (G.s) and P. aestuarii (P.a) co-cultures show two morphologically distinct species and their cell-cell contact via intimate extracellular associations (scale bars, 0.5 μm and 200 nm, respectively). These images are representative of the 20 SEM images and 26 TEM images obtained.
order to maintain the ΔombB-omaB-omcB-orfS-ombC-omaC-omcC mutant of G. sulfurreducens; however, it did not affect the growth of this strain of P. aestuarii, which we found to be naturally kanamycin-resistant (Supplementary Fig. 2b). Third, when the G. sulfurreducens mutant was genetically complemented with the ombB-omaB-omcB gene cluster, which was previously shown to restore extracellular electron transfer capacity to Fe(III) and electrodes (Fig. 4a), and then tested in the co-culture setup, light-dependent acetate consumption resumed (Fig. 4b).

*Geobacter* species require direct contact through outer membrane cytochromes and/or conductive pili for external electron transfer of electrons to insoluble electron acceptors (for example, electrode, Fe(III) oxide) and other organisms\(^ {23–25} \). Electron micrographs of the co-culture showed intimate connections between cells of G. sulfurreducens and P. aestuarii.

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**Figure 3** | Variation in acetate concentration in the co-culture of *P. aestuarii* and *G. sulfurreducens* in the presence and absence of light. (a) Switching the co-culture from a dark to an illuminated condition stimulated acetate consumption by the co-culture. (b) Switching from an illuminated to a dark condition stopped the photosynthesis of *P. aestuarii*, and therefore terminated acetate consumption by the co-culture. (c) Cell density (OD\(_{600}\)) of the co-culture of *P. aestuarii* and *G. sulfurreducens* in the presence and absence of light. The error bars represent the s.e.m. of replicated experiments (n = 3).

**Figure 4** | Current production and acetate consumption in co-culture by mutant and wild-type strains of *G. sulfurreducens*. (a) The ΔombB-omaB-omcB-orfS-ombC-omaC-omcC strain of *G. sulfurreducens* did not produce current on the electrode (+300 mVAg/AgCl). The current was partially restored when it was complemented with the ombB-omaB-omcB gene. Three biological replicates were performed for each strain, and typical data are presented. The arrow shows the point where the reactors were inoculated. (b) Acetate consumption of co-cultures of *P. aestuarii* and wild-type *G. sulfurreducens* (*G.s*-wt); of *P. aestuarii* and ΔombB-omaB-omcB-orfS-ombC-omaC-omcC *G. sulfurreducens* (*G.s*-mt), and of *P. aestuarii* and complemented ΔombB-omaB-omcB-orfS-ombC-omaC-omcC *G. sulfurreducens* (complemented *G.s*-mt) which was introduced with the ombB-omaB-omcB gene cluster. The error bars represent the s.e.m. of replicate experiments (n = 3).
Connections included cell-to-cell contacts between the naturally protruding prosthecae of the phototroph and the cell surface and cell appendages of *G. sulfurreducens*. Moreover, haem-stained cell preparations from the co-culture showed an abundance of haem-stained filamentous structures that connected the cells (Supplementary Fig. 6c,d). The presence of these structures suggests that electron transfer between cells of *G. sulfurreducens* and *P. aestuarii* is mediated by haem-containing proteins. Collectively, these results support a direct interspecies electron transfer mechanism for syntrophic anaerobic photosynthesis.

**Discussion**

Based upon these results, we propose a conceptual model for direct interspecies electron transfer between *G. sulfurreducens* and *P. aestuarii* in syntrophic anaerobic photosynthesis. Electrons originating from acetate are transferred from the heterotroph to the autotroph and used by the latter to fix CO$_2$ into cell material. The transfer benefits both organisms because it allows *G. sulfurreducens* to oxidize acetate as an electron donor and dispose of the electrons to an electron acceptor (the steps necessary to drive its bioenergetics), while simultaneously supplying the electrons needed by *P. aestuarii* to support photoautotrophic metabolism (Fig. 5). We demonstrated that *P. aestuarii* can obtain electrons from electrodes that were polarized at $-600$ mV$_{Ag/AgCl}$. *G. sulfurreducens* is well known for its ability to transfer metabolic electrons directly to extracellular electron acceptors including electrodes$_{17,21,23,24,26}$. Although the formal potential of the *G. sulfurreducens* biofilm grown on the surface of an electrode was shown to be around $-400$ mV$_{Ag/AgCl}$ (refs 17,26–28), some studies have suggested that this bacterium can adjust its redox activity to the potential of the electron acceptor$_{12,18,19,29}$. In addition, *P. aestuarii* is capable of reducing elemental sulfur$_{21}$, which is a known electron donor for *P. aestuarii*$_{14,20}$. Thus, it is likely that in the co-culture the potential of *G. sulfurreducens* was lowered sufficiently for it to serve as a donor for *P. aestuarii* photosynthesis.

We here demonstrated the growth of a photoautotroph supported by direct electron transfer from a heterotrophic partner. Several recent studies point to syntrophic interspecies electron transfers between various anaerobes as a major means of heterotrophic carbon metabolism in anoxic microbial habitats$_{5,4,8,30}$. Syntrophic anaerobic photosynthesis broadens this concept to include electron transfer from heterotrophs to phototrophs and reveals a previously unknown form of syntrophy that links anaerobic photosynthesis directly to anaerobic organic carbon metabolism. Moreover, although demonstrated here with the morphologically distinct green bacterium *Prosthecochloris*, we suspect that other species of green (and even purple) bacteria—organisms that are widely distributed in nature$_2$—may also establish direct interspecies electron transfer relationships with electrogenic heterotrophic partners.

From an ecological perspective, syntrophic anaerobic photosynthesis could represent an important form of carbon metabolism in the anoxic zones of poorly mixed freshwater lakes, where sulfide limitations restrict the activities of anoxygenic phototrophs and a shortage of inorganic electron acceptors limits anaerobic respiration. In addition to the ecological importance of direct electron transfer between heterotrophs and phototrophs, the discovery of syntrophic anaerobic photosynthesis reveals new possibilities for bioengineering phototrophic microbial communities for applications in the areas of waste treatment and bioenergy production.

**Methods**

**Bacterial strains and cultivation condition.** *Geobacter sulfurreducens* strain PCA (ATCC 51573) was used. The mutant strain of *G. sulfurreducens* in which the *ombB-omaB-omcB* gene cluster was replaced with an anthraquinone-2-sulfonate gene and its complement strain with the *ombB-omaB-omcB* gene cluster were created and characterized previously$_{22}$. *Prosthecochloris aestuarii* strain 728 was isolated from phototrophic mat harvested from epipsammal, hypersaline Hot Lake near Oroville, WA, USA$_{31}$. Strict anaerobic culturing procedures were used in serum vials sealed with butyl rubber stoppers throughout the study. *G. sulfurreducens* wild-type and mutant strains were cultured in a modified *Geobacter* medium containing acetate (10 mM) and fumarate (20 mM). The modified medium was determined to support the growth of *P. aestuarii* also when amended with sulfide and co-cultivation of both strains. The modified medium was made by dissolving 8.78 g of NaCl, 3.8 g of KCl, 0.25 g of NaHCO$_3$, 70 mg of CaCl$_2$, 2H$_2$O, 0.5 g of KH$_2$PO$_4$, 0.5 g of MOPS, 1.0 g of MgSO$_4$·7H$_2$O and 2.5 g of NaHCO$_3$ into 11 of medium to which were added 1× Wolfe’s mineral solution and 1× Wolfe’s vitamin solution. The compositions of these trace mineral and vitamin solutions were previously described$_{24}$. The medium was adjusted to pH 7.2, distributed into pressurized vials, boiled and purged with CO$_2$N$_2$, mixed gas (20%80%) before being capped and sterilized with an autoclave. Thioulsulfate (0.5 mM) was added for *P. aestuarii* biosynthesis to the media that were used in the cultivation of *P. aestuarii* on electrodes and in co-culture experiments. Kanamycin (200 μg ml$^{-1}$) was added to the media that were used for cultivation and co-culture experiments with *G. sulfurreducens* mutant strains$_{22}$. *P. aestuarii* was routinely cultivated anaerobically in sulfide-containing medium. Sodium sulfide (3.5 mM as Na$_2$S·9H$_2$O) was supplied to the *Geobacter* medium modified as described above. The sulfide-containing medium was allowed to sit in the dark for at least 2–3 h before being inoculated with *P. aestuarii*. The cultures were cultivated under constant illumination (8.3 ± 0.2 μmol m$^{-2}$ s$^{-1}$) and temperature (27 ± 0.4 °C) introduced by an incandescent light bulb (25 W). Co-cultures of *G. sulfurreducens* and *P. aestuarii* were obtained from pure cultures that were in their stationary phase. Before being used to inoculate co-cultures, the supernatant from each pure culture (16 ml) was removed after centrifugation (4,200 g, 20 min). Cell pellets were then washed with 2 ml of substrate-free medium without sulfur or sulfide. The co-cultures were initiated with a 0.5-ml inoculum of washed *G. sulfurreducens* and a 0.5-ml inoculum of washed *P. aestuarii* added to 15 ml of modified *Geobacter* medium containing 10 mM of acetate as the sole electron source. Oxygen scavengers such as sodium sulfide or cysteine were omitted from the media. The cultures were also incubated under constant illumination (8.3 ± 0.2 μmol m$^{-2}$ s$^{-1}$) and temperature (27 ± 0.4 °C) introduced by an incandescent light bulb (25 W). In some experiments, the co-culture vials were wrapped with aluminum foil to exclude light. The transferred co-cultures were initiated by inoculating fresh medium containing acetate as the sole electron donor with 1 ml of previously grown co-culture. The co-cultures were incubated under the same conditions as described above.

**Analytical methods.** To determine the change in cell density during culture, 1 ml of culture medium was sampled during incubation for optical density (OD) measurement using a UV–vis spectrophotometer (Agilent, CA, USA). The OD of *G. sulfurreducens* and co-cultures were determined at a wavelength of 600 nm, while the OD of *P. aestuarii* cultures were monitored at 720 nm. The concentration of organic acids was determined using high-performance liquid chromatography (HPLC) equipped with an Aminex NPX-87H column (Bio-Rad, CA, USA). Sulfuric acid (5 mM) was used as eluent at 0.6 ml min$^{-1}$. Organic acids were detected at 210 nm using a UV detector (Agilent).
Cell enumeration. Cells in the co-cultures were sampled at various times during growth. Samples of 1 ml were centrifuged (11,300 g, 10 min), and the supernatants were discarded. Cell pellets were washed three times with 0.1 M phosphate buffer (PBS; pH 7.2) before being fixed in 4% paraformaldehyde. Flow cytometry was performed using a BD Influx Fluorescence-Activated Cell Sorter (BD Biosciences, San Jose, CA, USA). To disrupt cell aggregates, the cultures were treated with 100 mM Na2EDTA (Sigma Aldrich), passed through a 25-gauge needle 25 times, and the samples were then fixed with SYBR Gold stain (ThermoFisher, Waltham, MA, USA). Optimization and calibration were performed before each flow cytometry analysis using 3.6-μm Ultra Rainbow fluorescent particles (Spherotech, Lake Forest, IL, USA). Forward and side scattering were used to gate out cellular debris, and the 488-nm argon laser was used to excite SYBR Gold while measuring emissions at 542/27 nm. Gating and median calculations for 20,000 cells were done using FlowJo software (Tree Star, Ashland, OR). The ratios of the two distinct populations of cells within a mixed microbial community were identified from 20,000 recorded cells using size and complexity gates within FCS Express (Los Angeles, CA, USA) flow cytometry software.

Electron microscopic analyses. Scanning electron micrographs (SEM) provided imaging of the cell morphology, attachment to electrode surfaces and distribution within co-culture biofilms. To sample co-culture biofilms of *G. sulfurreducens* and *P. aestuarii*, a glass piece or a membrane filter as the biofilm substrate was placed into the serum vials before inoculation. The glass piece or membrane filter with attached biofilms was carefully removed and fixed for SEM, as were the graphite electrodes with cells attached from bioelectrochemical systems (BESs). Samples were fixed in 2% paraformaldehyde, 0.1% paraformaldehyde in 0.1 M PBS and pH 7.2 for up to 12 h at 4 ºC, then washed three times in PBS for 10 min each, and dehydrated in ethanol series (10, 35, 50, 75, 95% and 3 × 100% for 10 min each). The samples were immediately immersed (twice for 10 min in hexamethyldisilazane (Sigma Aldrich, MO, USA) followed by air-drying for 9–12 h in a fume hood. The samples were sputter-coated with gold and imaged with a Quanta SEM (FEI, Hillsboro, OR) and Orion Helium ion microscope (Zeiss, Peabody, MA).

TEM whole mounts were prepared by applying a 5-μl drop of cells from a co-culture onto a formvar-coated grid (Electron Microscopy Sciences (EMS, Hatfield, PA), and air-dried. The grids were viewed under an FEI Tecnai TEM equipped with a 200-kV LaB6 electron source (FEI, Hillsboro, OR). The detection of haem using thin-section TEM was performed with 3.3’-diaminobenzidine (DAB, EMS) treatment, following the previously described protocols.26,30 Biocolloids from co-culture were collected by carefully removing medium. The biofilms at the bottom of a vial was transferred to centrifuge tubes, centrifuged at low speed (850 g, 5 min) and fixed in 2% gluteraldehyde, 2% paraformaldehyde in 0.1 M PBS overnight at 4 ºC. The fixative was replaced by three washes in PBS (10 min each) followed by two incubations (10 min each in the dark at room temperature) in PBS containing fresh DAB (0.05%). The haem stain was developed by the third incubation with PBS containing fresh DAB (0.05%) and 0.18% H2O2 (10 min in the dark at room temperature). DAB samples receive 0.7 mM DAB solution without H2O2. The reaction was stopped by washing three times in PBS before dehydration in an ascending series of ethanol and infiltration in LR White embedding resin (EMS) and cured at 55 ºC for 24 h. The polymerized blocks were sectioned to 70-μm thin sections with a Leica Ultracut UCT ultramicrotome using a Diatome (Switzerland) diamond knife. Ultrathin sections were then mounted on formvar-coated 100 mesh Cu grids and carbon coated. The TEM imaging was done using a Tecnai T-12 (FEI) with a LaB6 electron source, operating at 120 keV.

Construction of bioelectrochemical systems and operational conditions. The BESs for growing *P. aestuarii* were constructed as an H-type reactor consisting of two glass chambers (Adams & Chittenden Scientific Glass, CA, USA). The two chambers (100 ml working volume/each), which housed the counter and working electrodes, were separated by a cation-exchange membrane (5.07 cm2). Counter electrodes with cells attached from bioelectrochemical systems (BESs). Samples were filtered at 25 mm and immersed in electrolyte, creating a consistent projected surface of microbial community were identified from 20,000 recorded cells using size and complexity gates within FCS Express (Los Angeles, CA, USA) flow cytometry software.

Data availability. All relevant data are available from the authors upon request.

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Author contributions
P.T.H. designed and conducted the experiments and performed the analyses. I.S. created the Geobacter sulfurreducens mutant. M.T.M. isolated Prosthecochloris aestuarii from a microbial mat in Hot Lake (Washington, USA); A.C.D. conducted electron microscopy imaging. All authors contributed to experimental planning, data interpretation and writing of the manuscript.

Additional information
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