

Autotrophic and mixotrophic culture of electro-active microalgae with electron supplied from electrodes for CO₂ conversion

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Electrochemical technologies that involve microorganisms are considered to be promising for sustainable applications. Microalgae can be used for carbon capture through photosynthesis which can directly fix carbon dioxide (CO₂). The conversion of CO₂ into fuel energy and other high value metabolites without pollution can contribute to reduce CO₂ emissions with more economic value. Light energy to biomass conversion efficiency is a major challenge in microalgal cultivation. Electrode assisted cultivation techniques for improved photosynthetic and carbon (CO₂) fixation metabolism for growth and biomass productivity have rarely been explored for microalgae. Light limitation, which leads to the loss of photosynthetic efficiency that in turn leads to decreased microalgal growth, is a major problem in large scale cultivation systems. Here, we summarise the ability of microalgae to perform extracellular electron uptake from cathode material for efficient biomass production and CO₂ conversion. The present review provides insights into the possible development of electroactive microalgae under autotrophic and mixotrophic conditions for efficient CO₂ conversion. Using the current knowledge of bioelectrochemistry and learning lessons from electroactive bacteria, we propose a proof of concept for electroactive microalgae and their future applications in CO₂ sequestration.

Keywords: Electroactive microalgae, CO₂ fixation, Extracellular electron uptake, Autotrophy, Mixotrophy.

Introduction

Microalgae are unicellular photosynthetic organisms capable of producing high-value metabolites such as carbohydrates, lipids, proteins, polyunsaturated fatty acids, vitamins, and pigments. Microalgal biomass has garnered tremendous interest for producing nutraceuticals, pharmaceuticals, therapeutics, food supplements, feed, biofuel, bio fertilizers, etc. due to its high content of lipids and other high-value metabolites. Microalgal biomass has the potential to convert solar energy to organic materials and potential metabolites of nutraceutical and industrial value [1]. Microalgae have several advantages in comparison with higher plants. They have a high efficiency of fixing carbon dioxide (CO₂) and subsequently convert it into biomass and compounds of potential interest [2]. Ceramic carriers were used to achieve highest carbon reduction of approximately 53% by using photosynthetic bacteria [2]. Moreover, microalgae have a shorter life cycle and higher photosynthetic efficiency than higher plants. However, despite the research efforts made in the past few decades, the cost of microalgal products has

remained very high, compared with that of agricultural plant products. Hence, understanding the full potential of microalgae for the sustainable production of biomass and other products are necessary.

Electroactive microorganisms have gained substantial attention recently as they allow the flow and exchange of electrons between intracellular or extracellular redox electroactive donors or acceptors [3]. Electroactive microorganisms are currently applied in microbial fuel cells (MFCs) and microbial electrosynthesis (MES) [4]. Electro-activated microalgae are a proof of concept that the application of renewable electricity as an electron donor in microorganisms.

The application of renewable electricity in microorganisms like bacteria has already been exploited for the production of metabolites and biomass [5]. Currently, the focus of microbial electrosynthesis is limited to acetogens and methanogens [6]. Different types of mechanisms are involved in the transport of externally produced electrons to the electron transport chain (ETC) of host microorganisms [7]. Two types of electron transfer can be achieved: direct electron transfer (DET) and mediated electron transfer (MET). *Cupriavidus necator* shows MET using hydrogen as an electron mediator and presents a higher solar energy to biomass conversion rate of 9.7% compared with microalgae that have a solar energy to biomass conversion rate of 3% and 6% at large-scale outdoor

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cultivation systems and small-scale laboratory conditions, respectively [8, 9]. Due to ineffective light-to-biomass conversion, productivity typically decreases in photo bioreactors (PBRs), resulting in a lower photosynthetic efficiency (9-10%) than the theoretical maximum (corresponding to $\sim 80 \text{ g biomass m}^{-2} \text{ day}^{-1}$) [10]. Algal light conversion efficiency is reported to fall between 3% and 5% in PBRs used for large scale cultivation [11]. Therefore, the controlled cultivation of microalgae with efficient photosynthetic activity in a commercially viable manner is necessary.

Based on the physiology of microalgae, different cultivation systems and bioreactors can be developed to enhance light absorption and substrate utilization. Such systems should be able to provide sufficient nutrients and photons (not excess) to prolong the logarithmic growth of microalgae, which is necessary for sufficient product accumulation [12]. Mixotrophic cultivation systems can enhance growth and metabolite production in microalgae. Although these systems provide sufficient organic nutrients and energy for the growth of microalgae, they have a relatively poor photosynthetic efficiency ($\sim 2\text{-}6\%$) to fix CO_2 and only 5-10% of the allocated carbon is used to produce lipids and other secondary metabolites [13]. Therefore, metabolic and genetic engineering technologies are being considered to enhance microalgal production [14]. Among these, manipulation of the Calvin–Benson–Bassham (CBB) cycle and chloroplast electron transport chain are of potential interest for improving the photosynthetic efficiency of microalgae by increasing the efficiencies of CO_2 fixation and light harvesting [1].

Recently, external electron supplementation has gained increasing interest for enhancing the growth and metabolite synthesis in microalgae. Acceleration of carbon and energy flux in microalgae is associated with product biosynthesis. Lipid biosynthetic precursors like acetyl-CoA and NADPH are considered to be important factors for increasing product yield in microalgae [12]. For increasing the capture and delivery of electrons formed by substrate catabolism, extra energy is needed to accelerate the conversion of NADP^+ and NADPH in microalgae and this may produce less disturbance on the redox state than the strategy to increase the proportion of NADPH [1]. Therefore, providing electrons through external electrodes has the potential to accelerate energy flux and product yield in microalgae. Mixotrophic cultivation of microalgae using external electrodes can promote the ratio of NADPH/ATP in microalgal cells, which can boost CO_2 fixation and improve the yields of biofuels and high-value secondary metabolites from microalgae. Moreover, the external supplementation of electrons can prevent energy loss due to cell shading, which is the major reason for photosynthetic loss in microalgal cultivation systems.

Improving our knowledge of extracellular electron

uptake or electro-activated microalgae is critical from both biotechnological and industrial viewpoints [13]. Early research findings have proved extracellular electron uptake by autotrophic bacterial species [14]. Microalgae have been widely used as biophotovoltaics (BPV), wherein the transfer of electrons from microalgal cells to an external electrode takes place [15, 16]. Different types of photo electrodes are developed by researchers for this purpose [17, 18]. However, substantial research has not been conducted on the reverse flow of electrons. Owing to the industrial relevance of efficient microalgal cultivation systems and their emerging biotechnological applications, this review will focus on the possibility of extracellular electron uptake by microalgae in different cultivation modes to achieve maximum CO_2 fixation. In this review, we propose a possible mechanism of electro-activated microalgae to overcome the current limitations of CO_2 fixation and subsequent microalgal biomass production.

Challenges in microalgae cultivation

The industrial cultivation of microalgae is often carried out in open ponds or closed PBRs in the presence of sunlight and other chemicals to improve the biomass and metabolite accumulation [19]. When the microalgal cultures attain a high optical density, self-shading occurs, leading to an inhomogeneous distribution of light inside the cultivation system, which will limit the conversion of CO_2 . As a result, only the cells present at the top layers absorb sufficient incident light, leaving the bottom layers in light-limited conditions; this has major consequences on the photosynthetic metabolism of microalgae [20]. In light-limited conditions, cells do not have sufficient energy for the synthesis of NADPH/ATP, major factors for the Calvin–Benson pathway, and thus, the cells fail to sustain a high biomass growth rate. If the light intensity available is below the compensation point, energy is utilized for cell maintenance rather than CO_2 fixation [20]. Culture depth is an important parameter that affects the algal biomass production in large scale cultivation systems. Chisti reported that the culture depth suitable for high biomass production is up to 20 cm [21]. Even if cells are exposed to full sunlight, the chance of excess illumination is low. For example, photosynthesis in *Nannochloropsis* is reported to be saturated at a light intensity of $\sim 150 \mu\text{mol photons m}^{-2} \text{ s}^{-1}$, a value below the full sunlight intensity of $\sim 500\text{-}2000 \mu\text{mol photons m}^{-2} \text{ s}^{-1}$ (depending on the season) [22]. Microalgal cells can absorb sunlight efficiently even in saturated conditions, which produces 3Chl^* (triplet state chlorophyll) that cannot be used for photochemical reactions but generates reactive oxygen species (ROS). Generation of ROS molecules leads to the oxidation of proteins, lipids, and pigments with

photosystem II (PSII) D1 subunit as a major target [23]. The damaged proteins and other metabolites are subsequently degraded and resynthesized for regaining the photosynthetic activity [24]. The photosynthetic machinery of microalgae has been evolved to adapt to different light regimes to survive highly variable illumination under natural environments. Microalgae show non-photochemical quenching (NPQ) for the non-radiative de-excitation of excited chlorophyll molecules, which results in the dissipation of excess energy in the form of heat [20]. Antioxidant molecules are also involved in preventing photochemical damage to microalgal cells. Carotenoid molecules act as antioxidants as well as activators of NPQ in microalgal cells, with zeaxanthin being the major carotenoid molecule involved in this process. The cells synthesize zeaxanthin from violaxanthin by the activity of violaxanthin de-epoxidase enzyme under high light conditions; under light-limited conditions, zeaxanthin is converted back to violaxanthin by the activity of zeaxanthin epoxidase enzyme [25]. This photosynthetic regulatory mechanism plays a significant role in photon to biomass conversion efficiency. The repair of damaged PSII needs energy for the synthesis of proteins that can result in the decreased availability of nutrients for microalgal growth [10]. NPQ photo protection mechanisms dissipate energy as heat that decreases the photon to biomass conversion efficiency. For increased biomass productivity and photon to biomass conversion efficiency, an optimal balance should be maintained between photo protection and photon efficiency. In large scale cultivation systems, mixing for gas exchange can also affect light absorption and photosynthetic

efficiency. If the mixing is very fast, the zeaxanthin-mediated photo protection mechanism does not occur properly, leading to photo inhibition and decreased productivity [26].

Electron transport mechanism in microalgae

Microalgae are photoautotrophic organisms that utilize light as a primary energy source for their metabolism. Light is harvested through phycobilisomes that excites P700 and P680 chlorophyll pigments at the reaction centres of PS I and PS II (Photosystem I and Photosystem II) located in the thylakoid membrane of chloroplasts (Fig. 2). This process aids the extraction of electrons from water and releases oxygen as a by-product [27]. These electrons are then conveyed through PS II, cytochrome b6f, plastocyanin (PC)/Cyt c6, and PSI to ferredoxin (Fd). Ferredoxin acts as the distribution hub for photosynthetic electrons. The photosynthetic electron transport chain (PETC) generates a proton motive force (pmf) across the thylakoid membrane that acts as a driving force for the production of ATP from ADP and Pi by the enzyme ATP synthase (Fig. 1) [28].

Different types of fuel cells have been developed for wastewater treatment [29]. Electrosynthesis using photosynthetic microorganisms is an emerging field of research. Several studies have been conducted on phototrophic bacteria that have proved the underlying molecular mechanisms and flow of electrons from cathodes to the bacteria for energy transduction and biomass production [2, 6]. Most of the research on microalgae is focused on MFCs for electricity production

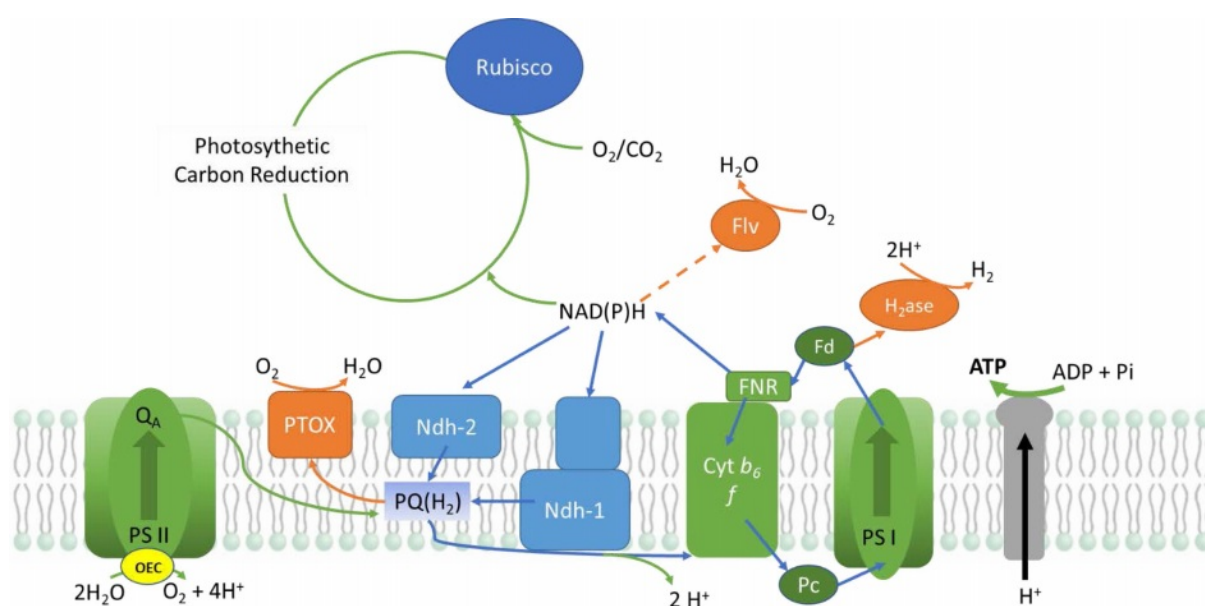


Fig. 1. Electron transport during light condition in microalgae (OEC; oxygen evolving complex, PSI & II; photosystem I & II, PTOX; plastoquinol terminal oxidase, PQ; plastoquinone, Ndh 1&2; NADH dehydrogenase like 1&2, Cytb6f; cytochrome b6f, PC; plastocyanin, FNR; ferredoxin-NADP reductase, Fd; ferredoxin, H₂ase; hydrogenase, Flv; flavin).

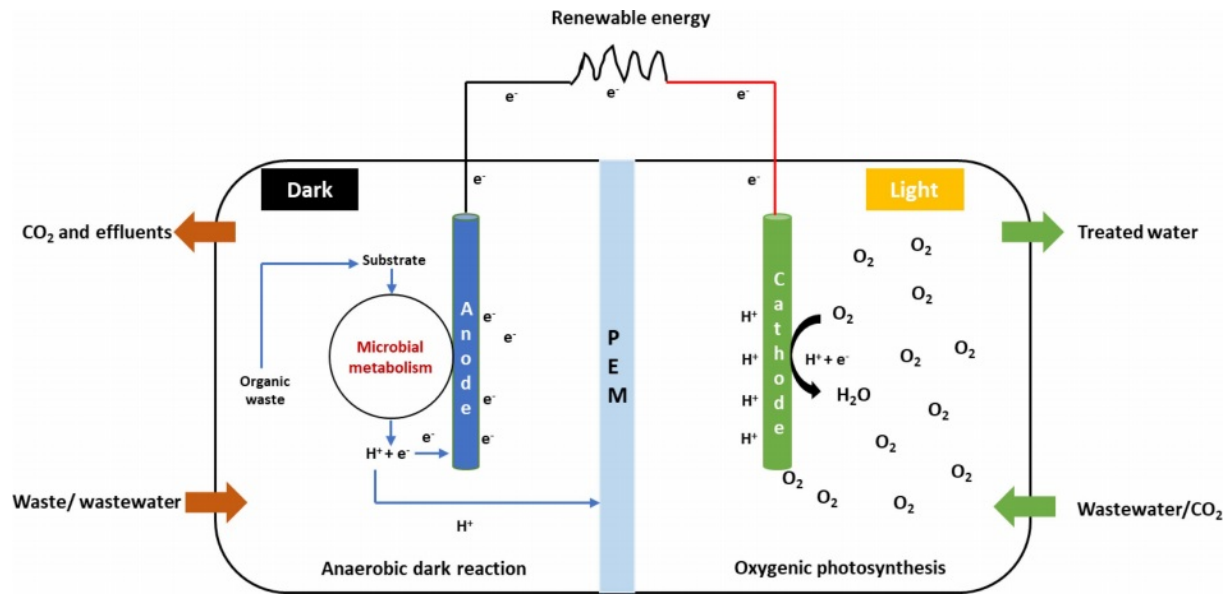


Fig. 2. Electron transport process in microbial fuel cell (MFC).

or wastewater treatment (Table 1). Microalgae that can accept electrons directly from cathode are referred to as electro-activated microalgae.

An MFC is a bio-electrochemical device capable of converting organic or inorganic substrates to energy through microbial metabolism [30]. Organisms capable

Table 1. List of microalgae used in MFC research and its applications.

Microalgae	MFC structure	Role of microalgae in MFC	Applications	References
Microcystis aeruginosa, Chlorella vulgaris	Two chamber	Substrate	Removal of trihalomethane	[76]
Arthrospira maxima	Two chamber	Substrate	Electricity generation	[77]
Chlamydomonas reinhardtii	Single chamber	Substrate	Electricity generation	[78]
Scenedesmus obliquus	Two chamber	Substrate	Electricity generation	[79; 80]
Chlorella vulgaris	Single chamber	Substrate	Electricity generation	[81]
Laminaria saccharina	Two chamber	Substrate	Electricity generation	[82]
Rhodobacter sphaeroides	Two chamber	Assisting anode	Solar powered MFC for electricity production	[83]
Chlorobium limicola	Two chamber	Assisting anode	Electricity generation coupled with dark metabolism	[84]
Rhodospseudomonas palustris	Single chamber	Assisting anode	Electricity generation	[85]
Chlorella vulgaris	Two chamber	Assisting cathode	Wastewater treatment and electricity generation	[34]
Chlorella vulgaris	Three chamber	Assisting cathode	Wastewater treatment and desalination	[86]
Chlorella vulgaris	Single chamber	Assisting cathode	Wastewater treatment and electricity generation	[87]
Microcystis aeruginosa	Two chamber	Assisting cathode	Electricity generation	[88]
Desmodesmus sp.	Two chamber	Assisting cathode	Electricity generation	[89]
Chlamydomonas reinhardtii and Pseudokirchneriella subcapitata	Two chamber	Assisting cathode	Wastewater treatment and bioenergy production	[35]
Synechococcus leopoliensis, Anabaena cylindrica and Chlorella pyrenoidosa	Two chamber	Assisting cathode	Electricity generation	[90]
Mixed culture of microalgae	Two chamber	Assisting cathode	Wastewater treatment	[90]

of doing this are known as electrogens or electro-chemically active microorganisms (EAM) that use the extracellular electron transport (EET) system for exchanging electrons with an insoluble material such as an electrode or metal oxide [31]. MFCs include the incorporation of microalgae in the cathode compartment that depends on the CO₂ produced at the anodic compartment as a carbon source. This process facilitates electrogenesis and can be used for wastewater treatment, CO₂ sequestration, and the synthesis of commercial products from microalgae (Fig. 2) [32]. Photo-cathodes with microalgae can be considered as a potential technique for the incorporation of photosynthesis into MFC systems [33] and have benefits such as nutrient removal, CO₂ fixation, supply of dissolved oxygen, and algal biomass production [34]. MFCs incorporated with microalgae are used for wastewater treatment for the removal of heavy metals and other nutrients, and the biomass produced by algae can be used to produce biofuels, biofertilizers, etc. [35, 36]. Commault et al. (2014) reported the use of photosynthesis in the cathodic compartment to replace the energy intensive mechanical aeration in bioreactors [37]. As a negative effect, the generation of a high oxygen content in the anodic compartment prevents electrogenesis even with high substrate degradation [38]. High electrogenic activity in the anodic chamber was observed by using photosynthetic bacteria as an anodic catalyst [18]. Oxygen availability at the cathode primarily depends on oxygenic photosynthesis with the help of PSI, PSII, and cytochrome b6f complex to transport electrons from water to NADP⁺. Small mobile molecules like plastoquinone and plastocyanin carry these electrons, thereby contributing to photosynthetic energy conversion [39].

Proposed mechanisms of electro-activation in microalgae

The carbon-derived chemicals produced from microalgae have been used in biofuel, nutraceutical, and pharmaceutical industries. However, the major energy source to produce these value-added commodities is photosynthesis, which provides the reducing power and energy. The addition of external electrons to photosynthetic electron transfer can effectively optimize biosynthetic pathways like the CBB cycle, thereby increasing the biomass and secondary metabolite production in microalgae. However, multiple factors should be taken into consideration for an effective mechanism of electro-activated microalgae.

Autotrophy, heterotrophy, mixotrophy, and photo-heterotrophy are the four major microalgal cultivation modes (Fig. 3). In the photoautotrophic mode of cultivation, microalgae utilize CO₂ as the carbon source and sunlight as the energy source for generating organic matter [40]. However, autotrophy is considered to be the most primitive way of microalgal cultivation and has been used since the 1950s in open pond system and closed PBRs [41].

In mixotrophy, the advantages of autotrophy and heterotrophy are combined to overcome the challenges of microalgal cultivation. In this mode of cultivation, microalgae can utilize both CO₂ and exogenous organic compounds such as glucose, fructose, acetate, and glycerol as the carbon source [42]. In mixotrophic cultivation, microalgae can grow heterotrophically using organic carbon for increased biomass and lipid accumulation and can also use inorganic carbon to produce oxygen through photosynthesis, thereby lowering the overall CO₂ emissions [43]. Unlike

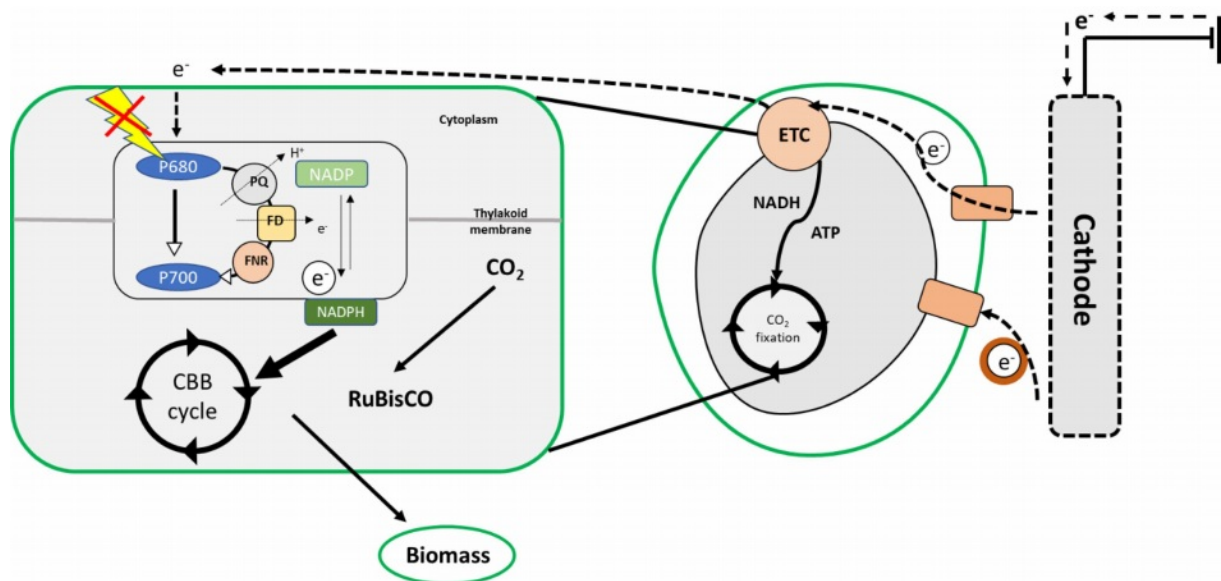


Fig. 3. Electron transport during electro-activation in microalgae.

heterotrophy, photosynthetic pigments are illuminated under the mixotrophic mode of cultivation [44]. Mixotrophic cultivation can be used to increase the growth and biomass production of microalgae with minimum contamination and a low cost [45]. The use of PBRs can reduce the chances of contamination even at an increased cost efficiency and a biomass production of ~5-15 g/L can be achieved, which is 3-30 times higher than that achieved under autotrophic cultivation conditions [46]. The mixotrophic cultivation of *Chlorella vulgaris* exhibited a biomass production of 2 g/L with glucose (1% w/w) as the carbon source [47]. Mixotrophic microalgal cultivation using glucose increased the growth rate of microalgae in comparison with photoautotrophic and heterotrophic cultivation methods [48]. Mixotrophic cultivation can be used to enhance lipid, carbohydrate, and protein accumulation in microalgae. Cultivation of *Asterarcys quadricellulare* showed a carbohydrate accumulation of 36.6% in media supplemented with 0.1 g/L [49]. Similarly, *C. vulgaris* produced a carbohydrate content of 8.74% using a mixture of glucose and glycerol as the carbon source [50]. The extracellular uptake of electrons by autotrophically and mixotrophically cultivated microalgae depends on various conditions like media composition, nature of electron mediators, and electrode material.

However, the phototrophic and mixotrophic cultivation of microalgae has many disadvantages in large-scale cultivation systems. The incorporation of electrons as an energy source in microalgal cultures can be considered to be a solution to this problem. In microbes, during extracellular electron transfer, cells try to make contact with each other through electrically conductive proteins, such as c-type cytochromes situated in outer membranes and Fe-S proteins, conductive pili, or periplasmic extensions, which aid the transfer of electrons across cell membranes [51]. The major technical difficulty in the use of electroactive microalgae is their thick cell wall and the membrane bound organelle chloroplasts, where photosynthesis occurs. This makes the eukaryotic algae different from cyanobacteria and other microorganisms owing to the difference in the relationship between primary electrogenic membranes and external electrodes [52].

Predicted extracellular electron transport proteins

The mechanism of PETC has remained stable during the course of evolution in cyanobacteria and algae except for the light harvesting complex proteins [27]. However, the extracellular electron transport proteins in cyanobacteria have been well studied by Nikkanen et al., 2021 [27]. They proposed the possible electron transport proteins in microalgae that are hypothetical proteins (Table 2). In cyanobacteria, NDH-1 acts as the major electron supplier to ETC followed by SDH,

cytochrome oxidase, PTOX, ARTO, FNR, etc. [53-55]. However, the presence of these proteins is predicted in the thylakoid membrane of different microalgal species that raises the possibility of extracellular electron uptake by eukaryotic microalgae under different cultivation conditions (Table 2).

Electron mediators

In photosynthetic microalgae, there is no evidence for direct extracellular electron transfer (DEET). However, bacteria like *Shewanella* show DEET ability with the help of nanowires situated in protrusions of the outer membrane [56]. In cyanobacteria, *Synechocystis* DEET has been reported with the help of conductive pili but the mechanism of DEET using pili is still unclear [57]. MET is widely employed in electroactive bacteria like methanogens and acetogens, and exploited using MFCs and bio-photovoltaic systems. Electron mediators have the ability to cross the semipermeable membranes of microalgae and interact with the inorganic or metallic electrode and the photosynthetic biological components [58]. The selection of a suitable electron mediator is necessary for the successful transfer of electrons from cathode to microalgae. The selected electron mediator should have the ability to pass the cell membranes through porin channels so that it can effectively transfer electrons within the cells [59]. Here, we propose a model wherein the anodic and cathodic compartments have the same electron mediator for transferring the electrons to the microalgae. For effective oxidation in the anodic compartment, the electron mediator should meet certain requirements. The mediator should not be toxic to the microalgae, and it should be effectively active for the electro-catalytic oxidation of various metabolites. Moreover, the electron mediator should be stable in the presence of different secondary metabolites and at different temperatures (10-40 C) and different pH ranges (5-9) [60]. There are several reports on the use of ferricyanide, quinone derivatives, neutral red, methyl viologen, etc. as electron mediators in MFCs to capture electrons from microalgae to generate bioelectricity [61]. However, in the case of electro-activated microalgae, the electron mediator should not be toxic to the algal cells. The currently used quinone derivatives are reported as algicides that are toxic to the microalgae at certain concentrations [62, 63]. Conversely, electron mediators such as riboflavin and Fe-EDTA are reported to promote the growth and electron transfer in biocathodes [64, 65]. The redox potential of an electron mediator is a significant factor that contributes to its activity. The redox potential of the artificial electron mediator should be positive enough to carry out fast reduction but negative enough to prevent energy loss. In addition, the kinetics of the microbial reduction and electrochemical oxidation should be as fast as possible [60].

Table 2. Predicted extracellular electron transport proteins in different microalgal species (Data obtained from UniProt, <https://www.uniprot.org/>).

Name	Function	Protein sequence ID
Plastid terminal ubiquinol oxidase 1	Terminal oxidase	>tr A8IU73 A8IU73_CHLRE Ubiquinol oxidase OS=Chlamydomonas reinhardtii OX=3055 GN=PTOX1 PE=3 SV=1
Plastid terminal ubiquinol oxidase 2	Terminal oxidase	>tr A8IEF7 A8IEF7_CHLRE Ubiquinol oxidase OS=Chlamydomonas reinhardtii OX=3055 GN=PTOX2 PE=3 SV=1
NADH Ubiquinone reductase	Reduces PQ and mediates cyclic electron transport without pumping additional protons	>tr A8JI60 A8JI60_CHLRE NADH:ubiquinone reductase (non-electrogenic) OS=Chlamydomonas reinhardtii OX=3055 GN=NDA2 PE=3 SV=1
Thylakoid membrane protein	Small extrinsic protein of thylakoid membrane, photosynthetic electron transport in photosystem I and response to high light intensity	>tr A8I547 A8I547_CHLRE Thylakoid membrane protein OS=Chlamydomonas reinhardtii OX=3055 GN=CHLRE_05g242400v5 PE=4 SV=1
Ferredoxin-NADP reductase	Key role in regulating the relative amounts of cyclic and non-cyclic electron flow to meet the demands of the plant for ATP and reducing power	>sp P53991 FENR_CHLRE Ferredoxin--NADP reductase, chloroplastic OS=Chlamydomonas reinhardtii OX=3055 GN=PETH PE=1 SV=1
Fe hydrogenase	Direct capture of photosynthetic electrons	>tr Q9FYU1 Q9FYU1_CHLRE Fe-hydrogenase OS=Chlamydomonas reinhardtii OX=3055 GN=hyd1 PE=1 SV=1
Light-harvesting complex stress-related protein 3.1	Nonphotochemical quenching, photosynthesis, light harvesting, protein-chromophore linkage, response to high light intensity, response to photooxidative stress	>sp P0DO19 LHR31_CHLRE Light-harvesting complex stress-related protein 3.1, chloroplastic OS=Chlamydomonas reinhardtii OX=3055 GN=LHCSR3.1 PE=1 SV=1
Photosystem II protein	Required for non-photochemical quenching (NPQ), a mechanism that converts and dissipates the harmful excess absorbed light energy into heat and protect the photosynthetic apparatus from photo-oxidative damage	>sp A8HPM2 PSBS1_CHLRE Photosystem II protein PSBS1 OS=Chlamydomonas reinhardtii OX=3055 GN=PSBS1 PE=1 SV=1
Uncharacterized protein Ycf 33	Electron transport	>tr A8J6Q3 A8J6Q3_CHLRE Uncharacterized protein ycf33 OS=Chlamydomonas reinhardtii OX=3055 GN=CHLREDRAFT_192514 PE=3 SV=1
Thioredoxin reductase	Electron transport	>tr A8HNQ7 A8HNQ7_CHLRE Thioredoxin reductase OS=Chlamydomonas reinhardtii OX=3055 GN=NTRC1 PE=3 SV=1
Flavodoxin-like domain-containing protein	Electron transfer activity	>tr A0A2K3CSH6 A0A2K3CSH6_CHLRE Flavodoxin-like domain-containing protein OS=Chlamydomonas reinhardtii OX=3055 GN=CHLRE_16g691800v5 PE=3 SV=1
Cytochrome c oxidase subunit 1	Cooperate to transfer electrons derived from NADH and succinate to molecular oxygen, creating an electrochemical gradient over the inner membrane that drives transmembrane transport and the ATP synthase.	>tr A0A650ANP2 A0A650ANP2_CHLVU Cytochrome c oxidase subunit 1 OS=Chlorella vulgaris OX=3077 GN=cox1 PE=3 SV=1
Cytochrome c oxidase subunit 2	Cooperate to transfer electrons derived from NADH and succinate to molecular oxygen, creating an electrochemical gradient over the inner membrane that drives transmembrane transport and the ATP synthase.	>tr Q6RFG7 Q6RFG7_CHLVU Cytochrome c oxidase subunit 2 (Fragment) OS=Chlorella vulgaris OX=3077 GN=Cox2 PE=3 SV=1
NADH ubiquinone oxidoreductase chain 4	Complex I functions in the transfer of electrons from NADH to the respiratory chain. The immediate electron acceptor for the enzyme is believed to be ubiquinone	>tr A0A650ANV6 A0A650ANV6_CHLVU NADH-ubiquinone oxidoreductase chain 4 OS=Chlorella vulgaris OX=3077 GN=nad4 PE=3 SV=1
Photosystem II protein D1	Electron transporter, transferring electrons within the cyclic electron transport pathway of photosynthesis activity	>sp P56318 PSBA_CHLVU Photosystem II protein D1 OS=Chlorella vulgaris OX=3077 GN=psbA PE=3 SV=1
Cytochrome b6	Component of the cytochrome b6-f complex, which mediates electron transfer between photosystem II (PSII) and photosystem I (PSI), cyclic electron flow around PSI, and state transitions	>sp P56321 CYB6_CHLVU Cytochrome b6 OS=Chlorella vulgaris OX=3077 GN=petB PE=3 SV=1

Table 2. Continued.

Name	Function	Protein sequence ID
Photosystem II protein D2	Electron transporter, transferring electrons within the cyclic electron transport pathway of photosynthesis activity	>sp P56319 PSBD_CHLVU Photosystem II D2 protein OS=Chlorella vulgaris OX=3077 GN=psbD PE=3 SV=1
Cytochrome b6	Component of the cytochrome b6-f complex, which mediates electron transfer between photosystem II (PSII) and photosystem I (PSI), cyclic electron flow around PSI, and state transitions	>sp P56321 CYB6_CHLVU Cytochrome b6 OS=Chlorella vulgaris OX=3077 GN=petB PE=3 SV=1
Thioredoxin reductase	Electron transport	>tr B9ZYY5 B9ZYY5_CHLVU Thioredoxin reductase OS=Chlorella vulgaris OX=3077 GN=CvNTR-C PE=2 SV=1
NADH-ubiquinone oxidoreductase chain 1	NADH dehydrogenase (ubiquinone) activity	>tr F1DGP3 F1DGP3_PHATR NADH-ubiquinone oxidoreductase chain 1 OS=Phaeodactylum tricornutum OX=2850 GN=nad1 PE=3 SV=1
Cytochrome c oxidase subunit 1	Cytochrome c oxidase activity	>tr F1DGN0 F1DGN0_PHATR Cytochrome c oxidase subunit 1 OS=Phaeodactylum tricornutum OX=2850 GN=cox1 PE=3 SV=1
Succinate dehydrogenase [ubiquinone] flavoprotein subunit	Responsible for transferring electrons from succinate to ubiquinone	>tr B5Y5N6 B5Y5N6_PHATC Succinate dehydrogenase [ubiquinone] flavoprotein subunit, mitochondrial OS=Phaeodactylum tricornutum (strain CCAP 1055/1) OX=556484 GN=SDH1 PE=3 SV=1
Predicted protein (similar functions to PGR5)	Photosynthetic electron transport in photosystem I and response to high light intensity	>tr B7FVH9 B7FVH9_PHATC Predicted protein OS=Phaeodactylum tricornutum (strain CCAP 1055/1) OX=556484 GN=PHATRDRRAFT_44748 PE=4 SV=1
Ferredoxin NADP (+) reductase	Ferredoxin-NADP+ reductase activity	>tr B7GCT8 B7GCT8_PHATC Ferredoxin--NADP(+) reductase OS=Phaeodactylum tricornutum (strain CCAP 1055/1) OX=556484 GN=PHATRDRRAFT_23717 PE=3 SV=1
NADH-ubiquinone oxidoreductase chain 4	NADH dehydrogenase (ubiquinone) activity	>tr A0A0U2F0A3 A0A0U2F0A3_BOTBR NADH-ubiquinone oxidoreductase chain 4 OS=Botryococcus braunii OX=38881 GN=nad4 PE=3 SV=1
Cytochrome c oxidase subunit 1	Cytochrome c oxidase activity	>tr A0A0U2F000 A0A0U2F000_BOTBR Cytochrome c oxidase subunit 1 OS=Botryococcus braunii OX=38881 GN=cox1 PE=3 SV=1
Cytochrome c oxidase subunit 1	Cytochrome c oxidase activity Electron transport coupled with proton transport	>tr A0A5B9R4G1 A0A5B9R4G1_HAELA Cytochrome c oxidase subunit 1 OS=Haematococcus lacustris OX=44745 GN=cox1 PE=3 SV=1
NADH-ubiquinone oxidoreductase chain 4	NADH dehydrogenase (ubiquinone) activity	>tr A0A5B9RCX0 A0A5B9RCX0_HAELA NADH-ubiquinone oxidoreductase chain 4 OS=Haematococcus lacustris OX=44745 GN=nad4 PE=3 SV=1
Ferredoxin--NADP reductase	Ferredoxin--NADP+ reductase activity	>tr A0A699ZA75 A0A699ZA75_HAELA Ferredoxin--NADP+ reductase (Fragment) OS=Haematococcus lacustris OX=44745 GN=HaLaN_16480 PE=4 SV=1
Uncharacterized protein Ycf33	Electron transport	>tr A0A699YIM3 A0A699YIM3_HAELA Uncharacterized protein ycf33 OS=Haematococcus lacustris OX=44745 GN=HaLaN_01554 PE=3 SV=1
Cytochrome c oxidase subunit 1	Cytochrome c oxidase activity Electron transport coupled with proton transport	>tr O47456 O47456_ISOGA Cytochrome c oxidase subunit 1 (Fragment) OS=Isochrysis galbana OX=37099 GN=COXI PE=3 SV=1
Succinate dehydrogenase	Electron transport	>tr A0A6T0Y6Z8 A0A6T0Y6Z8_ISOGA Succinate dehydrogenase (quinone) (Fragment) OS=Isochrysis galbana OX=37099 GN=IGAL00217_23638 PE=3 SV=1
NADH dehydrogenase [ubiquinone] iron-sulfur protein 4	Electron transport	>tr A0A6T1IPR9 A0A6T1IPR9_ISOGA NADH dehydrogenase [ubiquinone] iron-sulfur protein 4, mitochondrial (Fragment) OS=Isochrysis galbana OX=37099 GN=IGAL00217_50446 PE=3 SV=1
Ferredoxin--NADP(+) reductase	Ferredoxin--NADP(+) reductase activity	>tr A0A6T1AR05 A0A6T1AR05_ISOGA Ferredoxin--NADP(+) reductase (Fragment) OS=Isochrysis galbana OX=37099 GN=IGAL00217_30422 PE=3 SV=1

Cultivation conditions and reactor design

The cultivation of microalgae under different cultivation modes has various benefits and disadvantages. In phototrophic cultivation, light is the sole source of energy and when it comes to large scale cultivation systems, microalgae are not able to use light effectively. To solve this problem, mixotrophic cultivation systems have been introduced. However, in mixotrophy, the presence of expensive carbon sources and contamination with bacteria have been reported as major drawbacks. Thus, here we propose a mechanism to introduce external electrons as an additional source of energy in the normal growth medium under phototrophic or mixotrophic cultivation. The conductivity of the medium is a crucial factor for the cultivation of electro-activated microalgae. Moreover, the electron mediator and carbon source should work together for the efficient supply of electrons to the microalgal species selected.

Temperature, pH, applied current, and continuous nutrient supply are also crucial factors in microbial electrosynthesis [66]. The supply of gases such as CO₂ and light intensity also affect the MES in different bacterial and cyanobacterial species. However, reports on how these parameters affect microalgal electrosynthesis are still lacking. Changing the intrinsic properties of the reactor, such as temperature, salinity, and pressure, can clearly affect the viability of electroactive microalgae. Some researchers demonstrated the enhanced activity of MFCs under thermophilic temperatures above 45 °C that showed efficient substrate solubility, high microbial activity, efficient mass transfer, and low risk of contamination [67, 68]. However, most microalgae prefer mesophilic cultivation conditions and high temperatures may affect the viability of microalgae. The electrolyte nature and concentration can also affect electrosynthesis. The concentration of electrolyte significantly affects the conductivity of the cultivation medium and decreases the ohmic drop in the reactor [69]. Halophilic organisms have reported to have efficient MES activity [70]. This can be an advantage for marine microalgae that grow normally under halophilic conditions. The identification of marine microalgae with the ability of extracellular electron transfer and the modification of cultivation conditions could be effectively to develop electroactive organisms.

In addition, the electrolyte pressure can be used to enhance CO₂ availability in the microalgal culture. An optimal soluble CO₂ concentration is necessary for substrate specific consumption and maximum growth. The use of thicker electrode materials and different current densities can affect the available CO₂ in the bioreactor. Moreover, a high temperature and salinity have a negative impact on CO₂ availability. More research is needed to understand the factors that affect the electroactivity of microalgae under phototrophic and mixotrophic conditions. In addition, the optimization

of cultivation conditions and design of bioreactors are necessary.

Challenges and future perspectives

As a result of photosynthesis, microalgae can accumulate biomass and high-value metabolites. The application of an electric field has been shown to increase the growth and productivity of bacteria, fungi, and algae. The cellular machinery shows increased fermentation kinetics, increased synthesis of enzymes and RNAs, increased cell division, and increased mass transfer across the cellular membrane. The application of a low intensity electric field can significantly reduce the lag phase of *Saccharomyces cerevisiae* [71]. The polarization of *Volvox* sp. has been observed under the influence of electric stimuli [72]. The application of a static electric field of 2.7 kV/cm led to a 51% increase in the growth of *C. vulgaris* [73]. However, the application of electrostimulation on microalgal cultures for enhanced product and metabolite accumulation is still limited. The limited studies and finding an ideal candidate for electroactive microalgae are the current challenges in the development of a new production process.

Electro-technologies can be scaled up linearly in a cost-effective manner because they do not involve mechanical stress or diffusional processes. However, their industrialization faces several practical challenges like the requirement of a large electrode surface area, large electrode gap, large treatment chamber, and high flow rate. Moreover, effective electrode materials should be developed for successful microalgal cultivation in cathode compartment [74]. The need for generators increases at larger scales, consequently increasing the output voltage, high frequency pulses, and high energy capacity [75]. The major challenge in practical applications is finding power sources capable of producing high voltage and current outputs with a specific pulse and frequency. However, the development of generators and control systems for increased production in a cost-effective manner can solve the problem. More research is needed for the development of electro-activated microalgae to make them practically effective at a large scale.

Conclusions

The development of microalgae as a candidate for green chemical synthesis is currently challenging due to ineffective light to biomass conversion efficiency. The electroactive microorganisms have been gaining attention because of their high biomass and metabolite production rates. Electroactive microalgae are still in a proof-of-concept stage, and more research is needed to understand the external electroactive mechanism and production process. Advances in tools for screening

electroactive organisms can provide a path for the development of electroactive microalgae. It is also important to understand the rate-limiting factors in electroactive microalgae to scale up the production process. Future research in this field can make microalgal biorefinery more economically viable for chemical synthesis.

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