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# Cyanobacterial green chemistry: a blue-green approach for a sustainable environment, energy, and chemical production

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Increased human activity due to the ever-increasing global population has necessitated the urgent need for a sustainable environment, food, and energy. Cyanobacteria, classically known as blue-green algae, are oxygen-producing photosynthetic organisms that are emerging as an option to achieve sustainable development goals. These Gram-negative prokaryotes can efficiently sequester atmospheric CO2 due to an efficient carbon concentrating mechanism and divert it to the production of energy-rich compounds, i.e., biofuel, and other valuable chemicals, using their flexible metabolic chassis. Additionally, cyanobacteria also minimize the emission of methane, which is another greenhouse gas, by providing oxygen to methane-oxidizing bacteria. In recent years, several genetically engineered strains of cyanobacteria have been developed that can produce biofuels and several other valuable chemicals. Strains have also been engineered for bioplastic production and bioremediation purposes. These organisms have gained attention as biofertilizers and can increase the quality and fertility of soil. Thus, cyanobacteria are promising CO2 sinks that can contribute to global efforts in carbon capture and storage initiatives while producing bioenergy, cosmetics, pharmaceuticals, and several other valuable chemicals. Therefore, these blue-green cells can be used for green chemistry while minimizing the atmospheric CO2 concentration. In this review, we present various applications of cyanobacterial biomass to achieve sustainable development goals. We also discuss challenges associated with the wide application of cyanobacteria and the future direction to make full use of these robust organisms to fulfill our future demands in an environment-friendly manner.

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#### Sustainability spotlight

The sustainability of the environment, energy, and food is a major challenge associated with the continuous increase in global population. The greenhouse gas emissions due to fossil fuel burning and other industrial processes require a carbon-neutral society. The overexploitation of chemical fertilizers and other industrial contaminants is compromising the health of air, water, and soil. Also, the continuous piling of plastic waste, resistant to natural breakdown, is another environmental challenge. To overcome these challenges, greener ways of CO<sub>2</sub> sequestration, fertilizers, chemicals, and energy production are required without causing any negative impact on air, water, and soil. Cyanobacteria and microalgae are emerging biosystems that have the potential to fulfill future demands of sustainable food, energy, and the environment. In this work, we have discussed different features of cyanobacteria that can be used to sequester CO<sub>2</sub>, and sequestered CO<sub>2</sub> can be diverted through their versatile metabolic chassis using synthetic and molecular biology tools to produce food, energy, valuable chemicals, bioplastics, and water treatment systems. This work emphasizes the importance of the following UN sustainable development goals: zero hunger (SDG 2), good health and wellbeing (SDG 3), clean water and sanitation (SDG 6), affordable and clean energy (SDG 7), industry, innovation, and infrastructure (SDG 9), climate action (SDG 13), and life on land (SDG 15).

### 1 Introduction

The global population is continuously increasing and recently India has become the most populated country on the planet Earth. The ever-increasing population has both advantages and disadvantages. The advantage is to become a vast and growing

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consumer market to attract investment, increased workforce, and domestic production. However, along with these advantages, there are several disadvantages such as unemployment, poverty, scarcity of food, environmental degradation, overexploitation of natural resources, global warming, and environmental fluctuations, *i.e.*, climate change.<sup>2</sup> Climate change presents a multifaceted threat to global ecologies, societies, and economies and affects various environmental aspects such as extreme weather events, ice sheet melting, and sea level rise.<sup>3–5</sup> In response to the abovementioned environmental crisis, the

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Paris Agreement 2015 aimed to combat climate change and accelerate investments and actions to achieve a low-carbon sustainable future. Therefore, the rise in global temperature due to carbon dioxide (CO<sub>2</sub>) emissions and other anthropogenic gases underscores the urgent need for action to combat the negative effects of climate change.<sup>6</sup>

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In order to avoid damages from global warming and climate change, it is urgently required to limit greenhouse gas (GHG) emissions, particularly CO<sub>2</sub>, from various sources, in addition to sequestering it into biomass.<sup>7</sup> In the last two decades, cyanobacteria, classically known as blue-green algae due to their characteristic blue-green color, have emerged as promising organisms for the sustainability of the environment, agriculture, and energy.<sup>7-9</sup> Cyanobacteria play a crucial role in mitigating GHG emissions, mainly through CO<sub>2</sub> sequestration. As oxygenic photosynthetic Gram-negative prokaryotes, cyanobacteria utilize sunlight to convert CO<sub>2</sub> into organic compounds (Fig. 1), thereby controlling its concentration in the atmosphere.<sup>10,11</sup> Thus, cyanobacteria can act as a CO<sub>2</sub> sink due to the presence of a very effective carbon-concentrating mechanism (Fig. 1). These organisms accumulate inorganic carbon in the

form of bicarbonate inside the cells in a proteinaceous microcompartment called a carboxysome. 11-13

The special feature of efficiently sequestering atmospheric CO<sub>2</sub> makes cyanobacteria valuable allies in combating climate change and promoting environmental sustainability. In addition to CO<sub>2</sub>, cyanobacteria can also mitigate methane production, which is another GHG, *via* modulating microbial methane metabolism. Cyanobacteria promote the growth of methane oxidizing microorganisms (MOB) and utilize their metabolites. The physicochemical properties of soil are improved by cyanobacteria that can further promote methane mitigation by MOB.<sup>14</sup> In the context of climate change mitigation, cyanobacterial role as a potent CO<sub>2</sub> sink provides a valuable tool for reducing atmospheric carbon levels. The efficient photosynthetic conversion of CO<sub>2</sub> coupled with high biomass production by cyanobacteria makes these organisms a key player in biological carbon sequestration efforts.<sup>7,11,15</sup>

In the last few decades, efforts have been made to enhance the usage of cyanobacteria by using genetic manipulation and synthetic and metabolic engineering approaches.<sup>16</sup> Through genetic engineering, cyanobacteria produce various kinds of

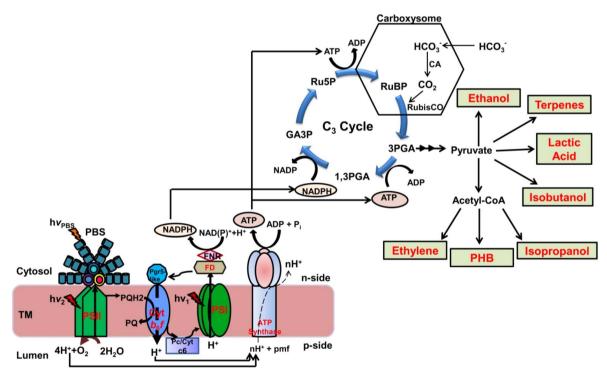


Fig. 1 Cyanobacterial biofactory for the production of valuable chemicals by sequestering atmospheric  $CO_2$ . The diagram depicts the integration of the photosynthetic light reactions and the Calvin–Benson–Bassham (CBB) cycle ( $C_3$  cycle) for the synthesis of various chemicals by diverting fixed carbon in cyanobacteria. Photosystem II (PSII) and photosystem I (PSI) generate a proton gradient across the thylakoid membrane in the lumen (p-side) and cytosol (n-side) using radiant energy. The proton gradient is utilized to generate ATP and NADPH that drive the  $C_3$  cycle in conjunction with the carboxysome. 3-phosphoglycerate (3PGA), which is produced in the carboxysome after fixation of  $CO_2$ , is converted into pyruvate. Pyruvate serves as a precursor for the synthesis of several chemicals, including ethanol, lactic acid, isobutanol, ethylene, and isopropanol. Additionally, acetyl-CoA derived from pyruvate is fed into the polyhydroxybutyrate (PHB) biosynthetic pathway to produce bioplastic, *i.e.*, PHB. For the synthesis of various chemicals using cyanobacterial metabolic chassis, the source of carbon is  $CO_2$  which is sequestered inside a proteinaceous microcompartment called a carboxysome in the form of bicarbonate (HCO $_3$ ). The whole process of sequestering  $CO_2$  and biosynthesis of various important chemicals by cyanobacterial metabolic chassis involves several enzymes. These enzymes can be inherently produced by cyanobacteria or can be installed into the cyanobacterial genome from distantly or closely related organisms using synthetic biology tools.

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Fig. 2 Cyanobacterial role in achieving green chemistry and environmental sustainability. Cyanobacterial cells use solar radiation, CO<sub>2</sub>, water, and nutrients to produce cell biomass that can be used for various purposes. Cyanobacteria can divert atmospheric CO2 to produce bioplastics, biofuels, biofertilizers, and other valuable chemicals. Cyanobacteria act as a CO2 sink and can assist in bioremediation and aerobic methane oxidation by methanotrophs.

chemicals, including alcohols, hydrocarbons, proteins, carbohydrates, carboxylic acids, isoprenes, toxins, antioxidants, pigments, vitamins, and methane. These chemicals have different applications in the biofuel, food, pharmaceutical, and cosmetic industries. Cyanobacteria are also useful in bioremediation and industrial biotechnology.16,17 Moreover, the use of cyanobacteria as biofertilizers offers a sustainable approach to enhancing crop yields by improving soil fertility and decreasing methane emissions in agricultural ecosystems such as rice fields. By leveraging the unique nitrogen-fixing abilities of cyanobacteria, farmers can enhance soil health, reduce the need for chemical fertilizers, and therefore, can contribute to environment-friendly agricultural practices. 18,19 These approaches not only benefit food security but also help in mitigating GHG emissions associated with conventional farming practices.

Cyanobacteria, with their unique capabilities and diverse applications particularly in environmental sustainability along with industrial biotechnology, hold significant promise for addressing critical global challenges such as climate change, food security, GHG emissions, and bioenergy production (Fig. 2). Thus, this review highlights the multifaceted role of cyanobacteria in achieving sustainability of the environment, energy, and chemical production while minimizing the levels of CO<sub>2</sub> and methane and fixing atmospheric dinitrogen.

## Cyanobacterial relevance in reducing greenhouse gas emissions

Cyanobacteria play a significant role in reducing GHG emissions through their unique metabolic processes. These photosynthetic microorganisms play a crucial role in carbon sequestration by absorbing atmospheric CO2 during photosynthesis, converting it into organic molecules and biomass (Fig. 1 and 2), and releasing oxygen as a byproduct. Additionally, several cyanobacteria are

capable of nitrogen fixation, and therefore, these organisms can enhance soil fertility and support plant growth while contributing to global carbon fixation.20 Also, cyanobacteria and their genetically engineered strains have been explored for their potential in biofuel production to offer a renewable and carbon-neutral alternative to fossil fuels.<sup>7,8</sup> By harnessing their capabilities in biofertilizers, methane reduction, carbon sequestration and fixation, and sustainable bioenergy production, cyanobacteria offer promising avenues for mitigating GHG emissions and addressing problems associated with global climate change (Fig. 2).

#### 2.1 Cyanobacterial oxygenation boosts methanotrophdriven methane oxidation in wetland rice cultivars

In present days, the discussion and research on methane production, release, and usage have gained significant attention due to its impact on global warming and atmospheric chemistry.21 Agricultural fields such as wetland rice fields, rivers, estuaries, oceans, wildfires, vegetation, and animals are identified as major natural sources of methane emission. In addition to natural sources, the majority of anthropogenic activities such as biomass burning and landfilling, animal rearing, and consumption of fossil fuels contribute to global methane emission.<sup>22</sup> Also, methane emissions in the atmosphere can increase as a result of population growth, waste production, and increased demand for food and fossil fuels. Nonetheless, methanogenesis in anaerobic flooded rice fields makes these agroecosystems one of the most important contributors to methane production.23

Agriculture is essential for both financial and food security, but it needs to be environment friendly. Therefore, to feed the increasing global population, the production of rice grains in the future will require the use of environment-friendly fertilizers and genetically engineered rice cultivars that can trap more carbon in the grains to minimize the emission of methane.<sup>24</sup> It is imperative to increase agricultural productivity without sacrificing the sustainability of the environment, and therefore, even a simple strategy can make a significant difference. According to Cuellar-Bermudez (2019),<sup>25</sup> cyanobacteria hold great promise for addressing the problem of global warming which is caused by GHG released due to natural and human activities. In addition to lowering the required quantity of nitrogen fertilizers, cyanobacteria can increase rice production with a simultaneous reduction in the amount of methane generated during rice cultivation.<sup>26</sup>

Thus, in the agricultural ecosystem, the applications of cyanobacteria can increase the production of grains, maintain the fertility and texture of the soil, and decrease methane yield. The methane produced by methanogenic bacteria in rice fields is utilized by methanotrophs which are obligate aerobic autotrophic microorganisms.<sup>27</sup> Methanotrophs convert methane into CO2 to obtain energy from its oxidation.28 The CO2 produced during the oxidation of methane can be consumed by cyanobacteria. Thus, methanotrophs play an important role in eliminating methane produced during rice cultivation or from any other sources. However, the oxidation of methane in waterlogged habitats is limited by the availability of oxygen.29 Therefore, inoculation of cyanobacteria and/or algae in waterlogged conditions, particularly in rice fields, can overcome the limitation of oxygen and promote the oxidation of methane by methanotrophs. In addition to promoting the oxidation of methane, the use of Anabaena sp., Nostoc sp., Calothrix sp., and Tolypothrix sp. as cyanobacteria-based biofertilizers enriches the soil quality and fertility by secreting several chemicals and fixing atmospheric nitrogen.7,30 However, excessive use of chemical fertilizers can limit the growth of microorganisms and, therefore, can negatively impact soil quality as well as consumption of methane by methanotrophs.

As a proof-of-concept, it has been shown that methane emission can be decreased by the use of cyanobacterial/algal biofertilizers due to the availability of oxygen produced during photosynthesis by these organisms to increase the oxidation of methane.<sup>30</sup> Thus, the integrated approach of using cyanobacterium-based biofertilizers not only improves the quality and fertility of the soil but also decreases the net production of methane from the agricultural field to attain sustainable agricultural practices.

# 2.2 Harnessing cyanobacteria as CO<sub>2</sub> sinks: an approach to climate sustainability

The primary cause of increasing  $\mathrm{CO}_2$  levels in the atmosphere is the burning of fossil fuels such as petroleum, coal, and natural gas which ultimately leads to global warming and climate change. Human activities, particularly industrialization and excessive fossil fuel burning mainly due to transportation, have significantly increased GHG emissions. To address the problems associated with global warming and climate change, various physical and biological methods have been employed to mitigate  $\mathrm{CO}_2$  levels.

Physical means of CO<sub>2</sub> sequestration have many disadvantages: having high costs associated with capturing,

transporting, and storing CO<sub>2</sub>. However, biological sequestration of CO<sub>2</sub> using cyanobacteria has been proposed to be a highly effective approach for decreasing the level of atmospheric CO<sub>2</sub>. <sup>7,33,34</sup> The CO<sub>2</sub> sequestered by cyanobacteria can be diverted for the production of valuable green chemicals, nitrogen or protein-rich biomass, and biofuels using flexible metabolic chassis of wild-type or genetically engineered strains (Fig. 1). <sup>7,8,15</sup> Cyanobacteria are selected primarily due to their efficient CO<sub>2</sub> sequestration ability and photosynthetic conversion of CO<sub>2</sub> to sugar, various industrially important bioactive molecules, and biomass production. <sup>8,11,15,35</sup>

Cyanobacteria are found in diverse habitats, including marine and freshwater ecosystems, and possess a carbonconcentrating mechanism (CCM) that enables them to accumulate a ~1000 times higher concentration of CO<sub>2</sub> than the ambient level. 11,12,35,36 The absorbed CO2 is accumulated around the Rubisco enzyme in a proteinaceous microcompartment called a carboxysome where its further reduction takes place to lock the sequestered CO2 into simple sugar. 15,37 The cyanobacterial CCM involves transporters for inorganic forms of carbon (Ci) such as CO2 and HCO3 along with carboxysomes that encapsulate Rubisco and carbonic anhydrase (CA) enzymes.11,12,38 It is believed that the cyanobacterial CCM evolved to minimize photorespiration that came into effect due to changes in the absolute and relative levels of CO2 and oxygen in the environment after the evolution of oxygenic photosynthesis.11,12

The cyanobacterial genome codes for C<sub>i</sub> transporters as single-gene products (e.g., BicA and SbtA) and some of the operons coding for transport machinery (e.g., BCT1, NDH-13, and NDH-14). Cyanobacteria have two CO2 absorption systems located in the thylakoid membrane and three transporters in the plasma membrane.11-13 Together, these five distinct inorganic carbon transporters make the CO2 acquisition system of cyanobacteria. However, all cyanobacteria may not have all five transporters that differ from each other in terms of their unique net affinity and uptake flux capacity for CO2 sequestration inside the cell.12,13 In order to mitigate the impact of growing CO<sub>2</sub> concentrations in the atmosphere, the potential of cyanobacteria to sequester CO2 due to the presence of an efficient CCM is getting recognition. Cyanobacteria play a significant role in the overall photosynthetic conversion of solar energy and CO2 assimilation which is almost ten to fifty times faster than in terrestrial plants.39 Therefore, using the potential of these biological systems is one of the most promising ways to lower the atmospheric CO2 level to mitigate the negative impacts of global warming.7 Currently, trees are a potential carbon capture solution in cities; however, their usage to capture CO<sub>2</sub> may be restricted due to their density, rate of growth, and availability of space.

Also, the high levels of atmospheric CO<sub>2</sub> together with other contaminants can promote stomatal closure to affect the uptake of CO<sub>2</sub> by plants.<sup>39</sup> The availability of green space is continuously decreasing in urban areas due to increasing population density, buildings, and other constructions. Therefore, an alternative approach must be taken to ensure that cities have a zero-carbon balance. The method of utilizing cyanobacteria/

algae to capture CO2 and other toxins from air, soil, and water into their biomass is called bio-capture or phyco-capture of CO2.7,40,41 As an alternative to large areas occupied by trees, the growth of cyanobacteria in photobioreactors has gained attention as an alternative approach to capturing carbon. 42

It is estimated that almost half of the oxygen available in the atmosphere is produced by cyanobacteria and microalgae which is released during the conversion of CO2 and solar radiation into chemical energy in the form of sugars. 15,36 According to Farrelly et al. 2013,43 these organisms can absorb up to 2.35 gigatons (Gt) of CO<sub>2</sub> per 100 000 km<sup>3</sup> culture area. It is also suggested that carbon fixation by microalgae could be a significant proportion of the remaining unidentified carbon sink.44 Thus, cyanobacteria together with other microalgae have the potential for CO<sub>2</sub> sequestration to tackle side-effects of global warming due to CO2 emission.

### Cyanobacteria can promote sustainable agricultural practices

The Green Revolution entered the global economy in 1965 and opened up new avenues for the agriculture sector which is now regarded as the backbone of the economy and provides several opportunities to a large portion of the population.45 While excessive use of chemical fertilizers has a positive effect on crop productivity, it could negatively impact the structure and physiochemical properties of the soil as well as the biotic and abiotic components of an ecosystem. These chemicals affect beneficial microbes along with nematodes and insects that are known to improve soil properties. 45,46 However, in the current scenario of continuous increase in global population, there is a need to further increase global food production without compromising the environment. Therefore, microbe-based biofertilizers present an excellent alternative to chemicalbased fertilizers for sustainable agricultural practices.19

Microbe-based fertilizers can reduce the usage of chemical fertilizers and increase crop production by improving the quality and fertility of soil. Among microbe-based fertilizers, several cyanobacteria are considered potent biofertilizers because of their ability to fix atmospheric nitrogen, secrete secondary metabolites, and improve soil fertility and structure.19,47 Cyanobacterial biomass enhances the physicochemical properties of deteriorated soil and increases soil water-holding capacity and mineral nutrient status.48 By naturally fixing atmospheric nitrogen, solubilizing phosphates, and producing chemicals that promote plant growth, biofertilizers supplement soil with additional nutrients. Biofertilizers are substances that hold microbial inoculants, artificially multiplied cultures of certain soil microbes that colonize the rhizosphere. However, by increasing the supplement or availability of primary micro- and macronutrients and growth-stimulating agents on the target crop, these microbial inoculants enhance soil fertility and crop productivity.49

There are approximately 1011 microbial cells per gram and more than 30 000 prokaryotic species found in the rhizosphere which is a narrow zone of soil that surrounds plant roots and

generally contributes to plant productivity. 49,50 In a natural agroecosystem, the interaction between bacteria and plants in the rhizosphere influences crop health by offering many services to crop plants such as decomposing organic matter, acquiring nutrients, absorbing water, recycling nutrients, and controlling weeds. 19,51 In addition to cyanobacteria, plant growthpromoting rhizobacteria (PGPR), phosphorus-solubilizers, sulfur-oxidizers, mycorrhiza, bacteria that can suppress diseases, endophytes that can withstand stress, and decomposers of organic matter are examples of other bioinoculants that function as biofertilizers.<sup>52</sup> Thus, cyanobacterial application in agricultural fields can promote sustainable agricultural practices by increasing soil quality and fertility without using chemical fertilizers.

### Cyanobacterial CO<sub>2</sub> capture: a green pathway to renewable energy rich chemical production

The whole world is looking towards options for renewable energy production to combat global climate change, limited availability of conventional energy resources, and their impact on GHG emissions. The recent G20 conference held in New Delhi witnessed the creation of the India-led Global Biofuels Alliance (GBA) in order to encourage the use of environmentfriendly biofuels. The aim of the GBA is to form a coalition between governments, international organizations, and business partners to promote the production and consumption of biofuels. It is estimated that global biofuel production needs to be increased three times to put the world energy system back on track with net zero carbon emission by 2050.7,53

The rise in biofuel demand is driven by several factors, including increasing oil prices, the urgent need to reduce GHG emissions from fossil fuel burning, the quest for energy independence among nations, and the opportunity for farmers to make a profit by providing feedstock and agricultural waste for biofuel production.54 Additionally, the production and utilization of biofuels are recognized as a carbon-neutral approach as they are produced from biomass that has sequestered a substantial amount of CO<sub>2</sub> from the atmosphere.<sup>7,55</sup> Globally, there is a growing trend in the production of biofuels from diverse bioresources through the application of innovative technology and biological processes. The utilization of agricultural crop waste and biomass to produce biofuels has been anticipated to lessen the negative impact of fossil fuel burning on the environment and the disposal of agricultural waste.7,56

Therefore, the production of biofuels from plants and microbial biomass has been the subject of investigation in the last few decades. 57,58 However, the production of biofuel from agricultural crops or waste can result in competition for fertile land to produce food and energy. To overcome this competition, additional research has been conducted on cutting-edge technology for the production of biofuels from alternative feedstock. Based on the type of feedstock used for the production of biofuels, there are four categories of biofuels (Fig. 3), i.e., first (1G), second (2G), third (3G), and fourth (4G) generations of

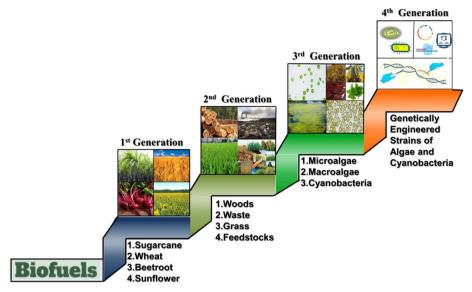


Fig. 3 Diagram depicting the evolution of different generations of biofuels. The first generation of biofuel uses crops such as sugarcane, wheat, beetroot, and sunflower, while the second generation biofuel utilizes wood, waste, and grass. The third generation of biofuel involves microalgae, macroalgae, and cyanobacteria, and the fourth generation employs genetically engineered strains of cyanobacteria and algae for enhanced production of biofuel.

biofuels.<sup>8,59</sup> A range of both gaseous and liquid biofuels such as biodiesel, ethanol, methanol, and methane are currently being produced from different biomass feedstocks.<sup>60-62</sup> Also, bioenergy produced from biomass is anticipated to contribute around 10–50% of the world's energy consumption by 2050.<sup>7</sup> Therefore, biomass-based energy production together with other sources of renewable energy such as solar and wind energy will play a significant role in fulfilling future global energy demand in a sustainable and environment-friendly manner.

The 1G biofuel is produced from edible food crops such as sugarcane, potato, oilseed, corn, barley, wheat, sunflower, and soybean.<sup>7,8</sup> Ethanol was the first energy-rich chemical produced from raw corn and sugarcane using fungal mycelia as a source for fermentation of sugars. 63,64 Fungi, e.g., Rhizopus sp. and Saccharomyces cerevisiae, can produce ethanol by fermentation of raw corn flour, and consequently, bioethanol has been produced on a large-scale from starch through enzymatic hydrolysis.65,66 The 2G biofuel is produced from cellulose and different organic waste materials such as wood, straw, and switchgrass and non-food plants such as Jatropha seeds. 7,8,67 3G biofuels are produced from eukaryotic algae and cyanobacterium feedstock, while the 4G biofuel production involves the use of genetically engineered strains of algae and cyanobacteria.<sup>7,8</sup> Genetically engineered strains having higher biomass production than their native strains or any other phenotypic alteration that gives them a competitive advantage over other strains are better suited for bioenergy production.

The quest for new energy alternatives has been further fueled by the urgent need to decrease the emission of CO<sub>2</sub> from fossil fuel burning. Therefore, genetically engineered strains of cyanobacteria and eukaryotic algae having higher CO<sub>2</sub>-sequestration efficiency can give an alternate option to energy sources while sequestering atmospheric CO<sub>2</sub>.<sup>11,68</sup> The algal or cyanobacterial biomass contains a large amount of carbohydrates, lipids, proteins, and several other constituents that can be used to produce different types of biofuels using physical and biological methods.<sup>7</sup> However, compared to algae and other photosynthetic organisms, cyanobacteria offer several advantages in biofuel production due to high growth rates leading to increased feedstock production, lower water, fertile land demand, simple and cost-effective nutritional requirements, and the ease of genetic manipulation through available trusted and tested genetic engineering tools.<sup>7,8,62</sup>

Biofuels derived from cyanobacteria offer a blend of alkanes, fatty acids, and fatty alcohols that closely resemble fossil fuels. This similarity of energy-rich molecules suggests that these biofuels could serve as excellent substitutes for transportation fuels without requiring significant alterations to vehicle engines.58 Over the past few decades, several attempts have been made to use photosynthetic organisms as living tools to convert solar energy into renewable energy. The production of ethanol by engineered Synechococcus elongatus PCC 7942, incorporating pyruvate decarboxylase and alcohol dehydrogenase II encoding genes from Zymomonas mobilis into its genome, marked a significant milestone in energy production using cyanobacteria. 69,70 Algenol Biotech, which is an industrial biotechnology company, established in 2009 in Fort Myers, FL, USA, uses genetically engineered cyanobacterial strains to produce bioethanol.

The genetic manipulation of basic cyanobacterial metabolic chassis has produced a number of compounds valuable for the energy sector (Fig. 1 and 4). The major advancement was made by diverting the carbon flux towards the production of ethanol in *Synechococcus elongatus* PCC 7942.<sup>69,71</sup> However, ethanol is not the greatest substitute for gasoline due to its lower energy

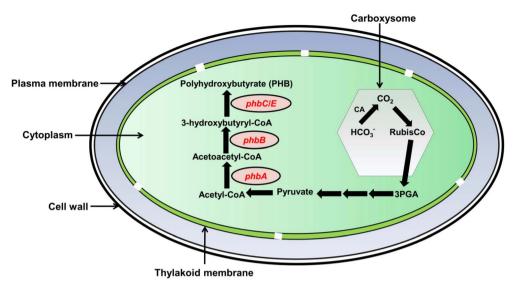


Fig. 4 Diagram illustrating the polyhydroxybutyrate (PHB) biosynthetic pathway within a cyanobacterial cell. The process begins with the conversion of pyruvate to acetyl-CoA. The enzyme encoded by the phbA gene converts acetyl-CoA to acetoacetyl-CoA, which is then transformed into 3-hydroxybutyryl-CoA by the phbB gene product. Finally, the product of phbC/E gene facilitates the polymerization of 3-hydroxybutyryl-CoA into PHB. The synthesis of PHB requires 3-phosphoglycerate (3PGA) as a source for its biosynthesis, and it comes out from the carboxysome after the reduction of  $CO_2$  by the RuBisCo enzyme. The carboxysome is the only microcompartment present in cyanobacteria and helps in the carbon concentrating mechanism to minimize the oxygenase activity of the RuBisCo enzyme.

content and ability to absorb water.72 Therefore, the focus has been shifted to developing energy-rich molecules with longer carbon chains. For example, isobutyraldehyde, which is a crucial ingredient for producing fuels from petroleum, has been produced in an efficient manner by modifying the valine biosynthetic pathway in S. elongatus PCC 7942.73 Despite these advantages, there are environmental concerns associated with the use of genetically modified (GM) cyanobacteria or any other microalgal strain for biofuel production. Potential issues include horizontal gene transfer and competition between GM strains and other microorganisms which could impact the natural balance of an ecosystem. However, large-scale cultivation of GM strains using closed photobioreactors can overcome the abovementioned challenges, but the economic viability of large-scale cultivation using expensive closed photobioreactors only for biofuel production still needs to be achieved.<sup>7,8</sup> Therefore, an integrated approach for the production of bioenergy and value-added molecules has been proposed to achieve economic viability.8 Also, the policy related to the use of genetically engineered strains of cyanobacteria and algae is still not clear at the global level.

# 5 Cyanobacteria in wastewater treatment: an eco-friendly approach for bioremediation

The global concern for the environment and human well-being has increased in the last few decades due to the addition of different pollutants to waterways.<sup>74</sup> Cyanobacteria exhibit versatile capabilities in addressing environmental issues, particularly in wastewater management and soil remediation.

These microorganisms contribute to removing excess levels of nitrogen and heavy metals from water systems and, therefore, have the ability to combat eutrophication. Cyanobacteria can metabolize complex compounds such as hydrocarbons and pesticides, and therefore, these organisms can be useful in promoting soil and water restoration.<sup>75-77</sup>

Domestic wastewater containing paper, human excrement, and synthetic detergent residues intermixes with industrial effluents containing a variety of organic and inorganic pollutants. Pesticides following their application permeate natural basins through soil erosion or atmospheric precipitation. Organic matter and suspended solids in wastewater impair light penetration and, therefore, hamper photosynthesis, promote sediment buildup, disrupt water self-purification processes, and diminish dissolved oxygen levels, whereas surface-active agents such as fatty acids and oils lower the oxygen content of water by producing a film layer. In the surface of the surfac

To address these issues, attention has been paid to biological wastewater treatment using cyanobacteria. <sup>79</sup> Cyanobacteria exhibit the exceptional capacity to decompose oil constituents and complex organics and sequester ions of zinc, cobalt, and copper metal. <sup>77,80</sup> Therefore, cyanobacteria exhibit promising solutions for treating secondary effluents derived from urban, agricultural, or industrial sectors. These microorganisms serve as powerful agents for the restoration and cleaning of diverse environmental settings as their distinctive metabolic characteristics and adaptive mechanisms allow them to effectively handle various pollutants. <sup>19,28</sup> In aquatic ecosystems, cyanobacteria excel in purifying contaminated water bodies by absorbing heavy metals such as mercury, lead, and cadmium, and therefore, these organisms can effectively minimize the hazardous consequences of these toxic substances in aquatic ecosystems. <sup>77,81,82</sup>

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However, optimization of cyanobacterial strains for specific pollutants, ensuring ecological safety, and scaling up bioremediation processes for large-scale applications still remain big challenges. To fully harness the potential of cyanobacteria, collaborative efforts involving genetic engineering and an enhanced understanding of their physiology are essential. This will permit the use of cyanobacteria in the ongoing fight against environmental degradation and conservation efforts. Cyanobacteria such as *Oscillatoria*, *Phormidium*, *Aphanocapsa*, and *Westiellopsis* have demonstrated their capability to eliminate nitrogen and phosphate ions from wastewater. Additionally, *Spirulina* strains possess diverse functional groups such as carboxyl, hydroxyl, and sulfate, which facilitate metal binding. Thus, the presence of these functional groups makes *Spirulina* strains effective for controlling pollutants such as zinc and nickel.

Also, research indicates that certain cyanobacteria such as Synechocystis sp., Westiellopsis prolifica, Nostoc hatei, and Anabaena sphaerica can break down organophosphorus or organochlorine insecticides in aquatic environments.84 However, widespread implementation of cyanobacteria in wastewater treatment is limited due to the efficient removal of biomass following treatment. Immobilization techniques using agarose, carrageenan, chitosan, alginate, and polyurethane foam can enhance the efficiency of nutrient or metal uptake with efficient removal of biomass from treated water.85 In natural ecosystems, several cyanobacterial species such as those belonging to genera such as Oscillatoria, Synechocystis, and Pleurocapsa form symbiotic associations with both aerobic and anaerobic microorganisms.84 Notably, cyanobacterial communities contribute to the breakdown of hydrocarbon pollutants found in crude oil spillover.85 While cyanobacteria do not directly decompose hydrocarbons, these organisms facilitate the process by supplying essential resources such as oxygen and nutrients to oil-degrading bacteria in a consortium.85

Therefore, to enhance the effective decomposition of hydrocarbons, an engineered consortium consisting of multiple cyanobacterial species, including *Phormidium*, *Oscillatoria*, and *Chroococcus* and the oil-degrading bacterium *Burkholderia cepacia* has been successfully developed and proven its efficiency for decomposing petroleum compounds.<sup>86</sup>

# 6 Polyethylene biodegradation: a sustainable approach to reducing plastic pollution

Polyethylene is a widely used material due to its affordability, durability, and versatility in various applications such as packaging, textile transportation, and manufacturing. However, its wide application is a major environmental challenge due to its improper disposal, especially in marine and urban environments.<sup>87</sup> Furthermore, its resistance to natural degradation is yet another environmental challenge. Due to the challenges encountered in the recycling of plastic waste, bioremediation gives an alternative approach to the breakdown of polyethylene by microorganisms.<sup>88</sup> In this process, microalgae and cyanobacteria can adhere to the surface of submerged polyethylene

and promote biodegradation by secreting substances that aid in the colonization of microorganisms. Some microorganisms can even directly use polyethylene as the sole carbon source. Divide bacteria and fungi have been extensively studied for polyethylene biodegradation, research on potential of cyanobacteria and other algae remains limited.

Notably, cyanobacteria and algae such as *Anabaena spiroides*, *Scenedesmus dimorphus*, and *Navicula pupula* have shown potential in degrading polyethylene. O According to Sarmah and Rout 2018, State two cyanobacterial species such as *Phormidium lucidum* and *Oscillatoria subbrevis* found on algal-covered polyethylene surfaces in sewage water are capable of utilizing polyethylene as a carbon source without any additional treatment. Thus, cyanobacteria together with microalgae offer a promising avenue for developing eco-friendly polyethylene degradation technology that is efficient, safe, and easy to implement. However, further intensive work is required to explore the possibility of using cyanobacteria and microalgae for the biodegradation of polyethylene.

# 7 Polyhydroxybutyrate (bioplastic) production and biodegradation

The degradation of polyethylene is extremely slow, and therefore, biodegradable polymers can be an alternative to a polyethylenefree environment. Polyhydroxybutyrate (PHB) is a biodegradable polymer that has gained significant attention due to its potential application as a sustainable alternative to conventional plastics in various industries.91 Cyanobacteria possess metabolic pathways that enable them to synthesize PHB as an intracellular carbon storage compound.92 PHB biosynthesis occurs via the condensation of acetyl-CoA molecules into D-3-hydroxybutyryl-CoA followed by polymerization and accumulation of PHB granules within cyanobacterial cells.93 Understanding the molecular mechanism and regulatory factors governing PHB biosynthesis in cyanobacteria is crucial for optimizing PHB production. Different cyanobacterial species such as Spirulina platensis, Gloeothece sp., Oscillatoria limosa, Synechococcus sp., and Synechocystis sp. accumulate considerable amounts of PHB.91

PHB is produced by three enzymatic reactions that start from acetyl coenzyme A (Fig. 1 and 4). Two acetyl-CoA molecules are condensed and converted to one acetoacetyl-CoA molecule by a 3ketothiolase enzyme encoded by the phbA gene.91 Furthermore, acetoacetyl-CoA is converted to D-3-hydroxybutyryl-CoA by NADPH-dependent acetoacetyl-CoA reductase which is encoded by the phbB gene. The final enzyme PHA synthase, encoded by phaC/phaE, forms PHB and also catalyzes the ester bond-forming process between the D-3-hydroxybutyryl moiety and an alreadyexisting PHB molecule.91 The enzymatic biodegradation of PHB by microorganisms such as algae, fungi, and bacteria involves a set of reactions that starts with the breaking of primary bonds of the polymer. This results in a change in the chemical structure of PHB and reduces its molar mass.92 The degradation of PHB begins with the action of PHA depolymerase, encoded by the phaZ gene, which acts on the PHB molecule and releases intracellular D-3-hydroxybutyrate. Furthermore, this molecule is

oxidized by the action of 3-hydroxybutyrate dehydrogenase to produce acetoacetate which is further esterified to acetoacetyl-CoA.  $^{92}$  Acetoacetyl-CoA is hydrolyzed by the *phaB* gene product to form acetyl-CoA which enters the tricarboxylic acid (TCA) cycle and ultimately degrades to  $CO_2$  and  $H_2O$ . The degradation of PHB produces  $CO_2$  and  $H_2O$  under aerobic conditions whereas  $CO_2$  and  $CO_3$  and  $CO_4$  are produced under anaerobic conditions.  $^{94}$ 

However, despite significant progress, several challenges remain to be addressed for the widespread commercialization of cyanobacterium-derived PHB.<sup>95</sup> These challenges include optimizing the rate of PHB production, improving the robustness and stability of cyanobacterial strains under dynamic environmental conditions, and reducing its production cost. Additionally, the development of scalable cultivation systems and downstream processing methods is essential for the industrial-scale production of PHB.<sup>96</sup> Therefore, future research efforts should focus on addressing the abovementioned challenges and exploring novel strategies for enhancing PHB production using cyanobacteria. Also, attention should be paid to screening new cyanobacterial isolates capable of producing PHB.

# 8 Cyanobacteria are biofactories for valuable chemical production (green chemistry)

The potential of genetically engineered cyanobacterial strains to directly convert CO<sub>2</sub> into target compounds provides an

environment-friendly and sustainable approach for green chemistry to produce various valuable chemicals (Fig. 1). Cyanobacteria offer an immense potential for genetic/metabolic engineering to produce a range of chemicals due to their unique features such as smaller genomes and genetic/metabolic diversity, rapid growth rates, and efficient mechanisms for CO<sub>2</sub> absorption.<sup>8,97</sup> Cyanobacteria are known to naturally produce a variety of enzymes and chemicals required for the functioning of their basic metabolic pathways. However, fine-tuning of gene expression or installation of novel metabolic pathways is crucial for the production of various chemicals on a large scale.<sup>8,98</sup>

Also, the production of various chemicals by cyanobacterial metabolic chassis can be designed in such a way that produced chemicals are secreted into the surrounding medium.<sup>99</sup> The development of cyanobacterial strains for the production of chemicals requires the integration of information obtained from basic physiological studies into modern-day studies such as biochemistry, molecular biology, synthetic biology, metabolic engineering, and systems biology. As depicted in Fig. 5, the developments in genetic tools, computational modeling, and high-throughput screening and sequencing techniques are continuously enhancing our capacity to engineer cyanobacteria for a variety of sustainable biotechnological purposes.<sup>100-102</sup>

Recent advancements in synthetic biology tools have facilitated the development of intricate metabolic engineering programs. A wide range of synthetic biology and metabolic

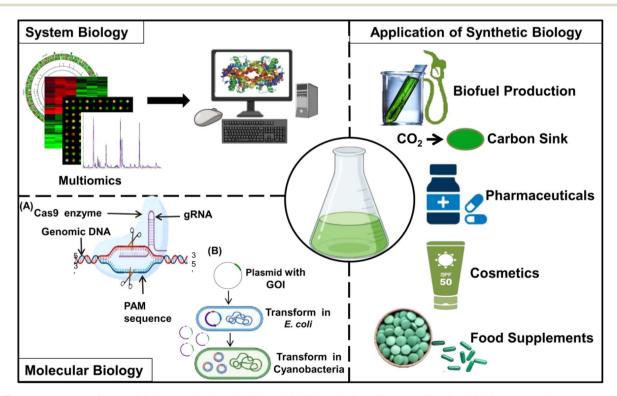


Fig. 5 The convergence of system biology and molecular biology highlights the broad impact of synthetic biology on various aspects of human welfare. System biology employs multiomics and computational analysis, while molecular biology utilizes genetic engineering tools such as CRISPR or classical methods of genetic modification in cyanobacteria for the development of improved strains. Together, these two approaches can help in developing new strains to sequester CO<sub>2</sub> and produce biofuels, pharmaceuticals, cosmetics, and food supplements.

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engineering tools have already been developed for cyanobacteria. These methods include utilization of the commonly used isopropyl-β-D-thiogalactopyranoside (IPTG)-controllable *trc* promoter, native and synthetic promoters, riboswitches, ribosome binding sites, selectable markers, suitable vectors for stable chromosome integration, dynamic regulation of gene expression, and genome-wide editing. 88,103,104

The availability of cyanobacterial genome sequencing data has enabled the integration of transcriptomics, proteomics, and metabolomics studies, leading to the development of genomescale models for various species such as Synechocystis sp. PCC 6803, Synechococcus sp. PCC 7002, Synechococcus elongatus PCC 7942, Anabaena sp. PCC 7120, and Arthrospira platensis NIES-39.105-107 Additionally, synthetic biology tools such as CRISPR/ Cas9-mediated gene editing have been successfully applied in cyanobacteria for precise modifications of the genome. 108,109 S. elongatus PCC 7942 was one of the first cyanobacteria, along with Synechocystis sp. PCC 6803, Anabaena sp. PCC 7120, and the fast-growing Synechococcus elongatus UTEX 2973 strain, where CRISPR editing has been successfully used. 108,109 However, CRISPR/Cas9-mediated gene editing in cyanobacteria, which is otherwise well established for other organisms, is still lagging behind due to the availability of more reliable and efficient homologous recombination-based approaches for genome editing using conjugation, natural transformation, or electroporation.

# 8.1 Green chemistry using genetically engineered strains of cyanobacteria

It is fascinating that *S. elongatus* PCC 7942 and *Synechocystis* sp. PCC 6803 have been successfully engineered for the production of energy-rich chemicals. As mentioned above, the production of ethanol by cyanobacteria was for the first time achieved by installing two genes of a Gram-negative anaerobic bacterium *Zymomonas mobilis*. Also, cellulose production in cyanobacterium *Synechococcus* sp. *leopoliensis* strain UTCC 100 through incorporation of the cellulose synthase encoding gene from *Gluconobacter xylinus* provides an alternative feedstock for ethanol production.

Another four-carbon alcohol, *i.e.*, butanol ( $C_4H_{10}O$ ), has been produced through microbial fermentation and from petrochemical propylene feedstock. Butanol and isobutanol are not naturally produced by cyanobacteria; however, genetic engineering has enabled the modification of cyanobacteria *Synechocystis* sp. PCC 6803 and *S. elongatus* PCC 7942 to produce these alcohols. PCC 6803 and S. elongatus PCC 7942 to produce these alcohols. The production of these energy-rich molecules by cyanobacteria has been achieved by introducing the genes from bacteria *Clostridium acetobutylicum*, *Treponema denticola*, and *E. coli*. 108,118,119

The production of isoprene, which is another valuable organic compound, has been accomplished by introducing the gene encoding isoprene synthase from the kudzu vine into *Synechocystis* sp. PCC 6803.<sup>120</sup> Likewise, the production of ethylene in *Synechococcus* sp. PCC 7942 and *Synechocystis* sp. PCC 6803 has been successfully achieved using their genetically engineered strains.<sup>121,122</sup> Once ethylene is produced by

cyanobacteria, a combination of biological and chemical conversion processes can be employed to transform ethylene into a range of hydrocarbons and functional groups that can be directly used as biofuels or as an intermediate for biofuel production. Thus, metabolic/genetic engineering offers potential means for utilizing renewable cyanobacterial feedstock to produce sustainable alternatives to conventional petroleum-based fuels using captured CO<sub>2</sub>.

In the cyanobacterium *Synechocystis* sp. PCC 6803, the production of isoprene was enhanced by introducing exogenous prokaryotic mevalonic acid (MVA) pathway genes.<sup>124</sup> Limonene, a cyclic monoterpene present in citrus fruit rinds, has various commercial uses in the flavoring and pharmaceutical industries. It is also a potential candidate for blending into jet fuel.<sup>125</sup> Limonene has been successfully produced using the cyanobacterium *Anabaena* sp. PCC 7120 by overexpressing key enzymes, *i.e.*, the limonene synthase gene from *Schizonepeta tenuifolia* and three endogenous terpene biosynthesis genes of *Anabaena* sp. PCC 7120.<sup>123</sup> Caffeic acid, a natural phenyl-propanoid with anticancer properties, was produced in the cyanobacterium *Synechocystis* sp. PCC 6803 by heterologous expression of a gene from *Arabidopsis thaliana*.<sup>126</sup>

The genetically engineered strains of cyanobacteria have also been used to produce various valuable compounds such as glycogen, mannitol, L-lactic acid, and D-lactic acid that are promising feedstocks for the medical, pharmaceutical, chemical, food, and biofuel industries. 127,128 Cyanobacteria naturally accumulate glycogen, which can be used as a feedstock for biofuel production. 129,130 Under specific conditions of high-light intensity, CO<sub>2</sub> concentration, nitrogen depletion, and moderate salinity, Synechococcus sp. PCC 7002 can produce 3.5 g per L glycogen with a productivity of 0.5 g per L per day. 131 The same organism has also been engineered to produce mannitol from CO2 by overexpressing mannitol-1-phosphate dehydrogenase (MtlD) and mannitol-1-phosphatase (Mlp) encoding genes of E. coli and Eimeria tenella, respectively. The engineered strain can produce 1.1 g  $L^{-1}$  of mannitol with a productivity of 0.15 g per L per day.132

L-Lactic acid used in various industries can be produced from pyruvate using an NADH-dependent lactate dehydrogenase (Ldh) from *Lactococcus lactis*. <sup>133</sup> The optimization strategies such as increased NADH-dependent lactate dehydrogenase (*ldh*) gene dosage and overexpression of the *Enterococcus faecalis* pyruvate kinase gene in *Synechocystis* sp. PCC 6803 has resulted in 840 mg per L L-lactic acid with a productivity of 22 mg per g dry cell weight per h. <sup>134-136</sup> D-Lactic acid, which is valuable for biodegradable plastics, can be produced by overexpressing the D-lactate dehydrogenase (D-Ldh) encoding gene. Overexpression of native and heterologous D-Ldh from *Lactobacillus delbrueckii* and inactivation of native poly-3-hydroxybutyrate and acetate pathways have resulted in an increased production of D-lactic acid in *Synechocystis* sp. PCC 6803. <sup>137,138</sup>

The production of 3-hydroxypropionic acid (3HP) and poly-3-hydroxybutyrate-*co*-4-hydroxybutyrate (P3HB4HB) has also been achieved in engineered strains of cyanobacteria *S. elongatus* PCC 7942 and *Synechocystis* sp. PCC 6803. In *S. elongatus* PCC 7942, two pathways, namely malonyl-CoA and β-alanine

dependent pathways, for 3HP production can be engineered.<sup>139</sup> However, in *Synechocystis* sp. PCC 6803, a different pathway has been engineered to directly reduce malonyl-CoA into 3HP by the overexpression of the malonyl-CoA reductase encoding gene of *Chloroflexus aurantiacus*.<sup>140</sup> In *S. elongatus* PCC 7002, P3HB4HB production has been achieved by expressing a heterologous poly-3HB synthesis operon (*phaABEC* operon of *Chlorogloeopsis fritschii* PCC 9212). The genetically engineered strain gives the production of approximately 4.5% P3HB4HB, containing 12% 4-hydroxybutyrate (4HB), of total dry cell weight.<sup>141</sup> Thus, genetic modification of wild-type cyanobacterial strains demonstrates the potential of these prokaryotes to be used as a host to produce biodegradable polymers using atmospheric CO<sub>2</sub>.

Cyanobacteria produce a variety of secondary metabolites, including peptides, alkaloids, and polyketides, exhibiting a wide range of biological activities.8 These metabolites are potential pharmaceuticals that can address the need for new drugs amid increasing antibiotic resistance and emerging diseases. Cyanobacterial peptides such as microcystins, nodularins, cylindrospermopsin, and cyanopeptolins show potent inhibitory activity against specific protein phosphatases and serine proteases.8 Therefore, these chemicals pose significant health risks to humans and animals. However, their specificity offers the potential for developing targeted therapies with fewer side effects in comparison to traditional drugs. These metabolites also contribute significantly to scientific research, particularly in understanding cell signaling and cancer biology. 142,143 Thus, cyanobacterial toxins are valuable candidates for drug development and hold considerable potential for biotechnological applications in medicine and agriculture.

Other cyanobacterial compounds exhibit diverse biological activities such as antibacterial, antifungal, antimalarial, antiprotozoal, and antiviral. Also, a few cyanobacteria show antitumor activity and antimitotic activity by inhibiting microtubule assembly and colchicine binding to tubulin. Thus, cyanobacterial metabolites can be used to develop new pharmaceuticals such as anticancerous agents to address some of the most challenging health issues of the present day. The ongoing research and discovery of new compounds from cyanobacteria could further lead to the development of novel therapeutics with targeted mechanisms of action. Also, the continued exploration and characterization of cyanobacterial metabolites will be crucial in harnessing their full potential for medical advancement in the future.

Cyanobacterial toxins also have commercial applications as algaecides, herbicides, and insecticides. For example, the indole alkaloid hapalindole A exhibits antialgal and antimycotic properties, while norharmane from *Nodularia harveyana* shows anticyanobacterial activity. <sup>146</sup> Cyanobacterin, produced by *Scytonema* sp., is a potent PSII inhibitor, and it is effective against algal and cyanobacterial blooms at low concentrations. <sup>147</sup> Similarly, cyanobacterins LU-1 and LU-2 from *Nostoc linckia* inhibit PSII-dependent electron transport. Other notable secondary metabolites that are effective against algal blooms are nostocyclamide, nostocine A, and nostocarboline from *Nostoc* species. <sup>148</sup> Additionally, pentacyclic calothrixins from

*Calothrix* sp. inhibit transcription and DNA synthesis, while microcystin from *Microcystis aeruginosa* acts as a growth inhibitor for several aquatic plants. Also, cyanobacteria are rich sources of photoprotective compounds, mycosporine-like amino acids (MAAs) and scytonemins. These compounds are crucial for their application in the pharmaceutical, cosmetic, and toiletries industries due to their stability under different photophysical and photochemical conditions.

In summary, while the primary focus on cyanobacterial toxins has been on their detrimental effects, there is growing interest in their potential advantages. These include their application in medical research and their ecological significance in aquatic systems. Therefore, further understanding the dual aspects of cyanobacterial toxins could lead to innovative approaches in both environmental management and biomedical science. This will also help in transforming a natural hazard into a resource for medical and environmental advancement.

### 9 Conclusions and future directions

The potential of cyanobacterial metabolic chassis to sequester CO<sub>2</sub> and convert it into valuable chemicals is immense, and genetically engineered strains can be designed for direct secretion of synthesized chemicals into the medium. For instance, as a proof-of-concept, Ducat et al. (2012)150 developed an engineered strain of S. elongatus PCC 7942 that can produce and secrete sucrose into the growth medium. The produced sucrose can be directly harvested from the liquid medium without the need for cell harvesting and downstream processing, or it can also be used to establish a consortium between cyanobacteria and heterotrophic organisms such as yeast or eubacteria. Thus, a consortium between cyanobacteria and heterotrophic organisms opens doors for sustainable biotechnological solutions where heterotrophic organisms can thrive without needing an external carbon source. However, the success of such a consortium relies on harnessing cyanobacterial metabolic networks to either replace or enhance current production systems for specific products. Fortunately, advancements in bioprospecting and the development of platform technologies have significantly expanded our understanding of cyanobacterial metabolism to enable precise manipulation and optimization for desired outcomes. These studies underscore the potential for cyanobacteria to play a pivotal role in sustainable bioproduction and, therefore, offer commercially viable alternatives to traditional methods.

One key advantage of cyanobacteria lies in their highly efficient CCM that enables them to effectively utilize bicarbonate and overcome the limitations posed by the slow diffusion of CO<sub>2</sub> into water. By enhancing the efficiency of the cyanobacterial CCM, future work should aim to increase the concentration of CO<sub>2</sub> around the enzyme Rubisco to improve photosynthetic efficiency and reduction of photorespiration in cyanobacterial strains targeted for chemical production. For example, an extra bicarbonate transporter has been installed in *Synechocystis* PCC 6803 which leads to a 2-fold enhancement in the growth rate and a higher biomass accumulation. <sup>151</sup> Continued research into optimizing the cyanobacterial CCM and exploring novel

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applications in carbon-capture technologies will be crucial in combating climate change and achieving a more sustainable future. By further exploring their diverse applications, advancement in genetic engineering techniques, and scaling up their production processes, cyanobacteria can play a pivotal role in shaping a more sustainable and environment-conscious future. Therefore, continued interdisciplinary research efforts and collaborations are essential in unlocking the full potential of cyanobacteria to address global challenges and foster a greener tomorrow.

The advancements in understanding and manipulating cyanobacterial metabolism coupled with the development of new platform technologies have significantly expanded our ability to harness these organisms for sustainable bioproduction. By combining the right tools with the right evanobacterial strain, we can achieve viable productivity and pave the way for the commercialization of cyanobacterium-based biotechnological solutions. Looking ahead, the future of cyanobacterium research and application presents exciting opportunities for innovation and impact. By harnessing the genetic and metabolic potential of these microorganisms, researchers can further enhance biofuel and food production, CO<sub>2</sub> sequestration, and specific chemical production to achieve sustainable developmental goals in a greener way. In conclusion, cyanobacteria have immense potential to tackle global concerns related to the sustainability of food, energy, and the environment; however, the economic viability of such attempts poses a major challenge. Therefore, governments need to subsidize the costs associated with cyanobacterial and algal cultivation to achieve a carbon-neutral environment.

### Data availability

No primary research results, software or code have been included and no new data were generated or analysed as part of this review.

#### Author contributions

PP, DP, and SPS conceptualized the idea. PP, DP, AG, RG, ST, and SPS wrote the manuscript. RG and ST prepared figures. SPS edited the manuscript. All authors reviewed the manuscript.

#### Conflicts of interest

The authors declare no conflicts of interest.

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### References

- 1 United Nations Population Fund (UNFPA), State of World Population Report, 2024, https://www.unfpa.org.
- 2 United Nations Population Fund (UNFPA), State of World Population Report, 2023, https://www.unfpa.org.
- 3 Intergovernmental Panel on Climate Change (IPCC), Climate Change 2022: Impacts, Adaptation and Vulnerability, 2022, https://www.ipcc.ch.
- 4 B. C. O'neill, M. Oppenheimer, R. Warren, S. Hallegatte, R. E. Kopp, H. O. Pörtner, R. Scholes, J. Birkmann, W. Foden, R. Licker and G. Yohe, *Nat. Clim. Change*, 2017, 7, 28–37.
- 5 M. Barnett, Manage. Sci., 2023, 69, 7562-7584.
- 6 Intergovernmental Panel on Climate Change (IPCC), Climate Change 2023: Synthesis Report, 2023, https://www.ipcc.ch.
- 7 P. K. Maurya, S. Mondal, V. Kumar and S. P. Singh, *Environ. Sci. Pollut. Res. Int.*, 2021, 28, 49327–49342.
- 8 R. Rajneesh, S. P. Singh, J. Pathak and R. P. Sinha, *Renewable Sustainable Energy Rev.*, 2017, **69**, 578–595.
- 9 J. Pathak, H. Ahmed, S. P. Singh, D.-P. Häder and R. P. Sinha, in *Aquatic Ecosystems in a Changing Climate*, CRC Press, 2018, pp. 45–61.
- 10 J. Zhou, H. Meng, W. Zhang and Y. Li, *Adv. Exp. Med. Biol.*, 2018, **1080**, 97–116.
- 11 P. Pandey, R. Gupta, S. Tiwari, A. Gupta, S. Mondal, R. P. Sinha and S. P. Singh, in *Cyanobacteria: Metabolism to Molecules*, Academic Press, 2024, pp 57–67.
- 12 G. D. Price, M. R. Badger, F. J. Woodger and B. M. Long, *J. Exp. Bot.*, 2008, **59**, 1441–1461.
- 13 M. R. Badger and G. D. Price, J. Exp. Bot., 2003, 54, 609-622.
- 14 P. Agarwal, R. Soni, P. Kaur, A. Madan, R. Mishra, J. Pandey, S. Singh and G. Singh, Front. Microbiol., 2022, 13, 939347.
- 15 S. Lucius and M. Hagemann, Front. Plant Sci., 2024, 15, 1417680.
- 16 S. Mazard, A. Penesyan, M. Ostrowski, I. T. Paulsen and S. Egan, Mar. Drugs, 2016, 14, 97.
- 17 D. C. Ducat, J. C. Way and P. A. Silver, *Trends Biotechnol.*, 2011, **29**, 95–103.
- 18 M. S. Massey and J. G. Davis, Nitrogen, 2023, 4, 253-262.
- 19 J. Pathak, R. Rajneesh, P. K. Maurya, S. P. Singh, D.-P. Häder and R. P. Sinha, Front. Environ. Sci., 2018, 6, 7.
- 20 L. Curatti, M. D. Nascimento, L. A. Pagnussat, L. Sanchez Rizza, A. O. Sanchez, L. Garcia Martinez and J. A. Hernandez, Rev. Environ. Sci. Biotechnol., 2024, 23, 291–320.
- 21 K. Senthilraja, S. Venkatesan, D. Udhaya Nandhini, M. Dhasarathan, B. Prabha, K. Boomiraj, S. Mohan Kumar, K. Bhuvaneswari, M. Raveendran and V. Geethalakshmi, Agriculture, 2023, 13, 1037.

22 R. J. Cicerone and R. S. Oremland, *Global Biogeochem. Cycles*, 1988, 2, 299–327.

- 23 J. Hu, M. Bettembourg, L. Xue, R. Hu, A. Schnürer, C. Sun, Y. Jin and J. F. Sundström, Sci. Total Environ., 2024, 920, 170980.
- 24 P. L. Bodelier, Nature, 2015, 523, 534-535.

Critical Review

- 25 S. P. Cuellar-Bermudez, J. A. Magdalena, K. Muylaert and C. Gonzalez-Fernandez, *Algal Res.*, 2019, 44, 101689.
- 26 R. Ambrosio, L. S. Rizza, M. D. Nascimento, H. G. J. Pacheco, L. M. M. Ramos, J. A. Hernandez and L. Curatti, in *Cyanobacterial Lifestyle and its Applications in Biotechnology*, Academic Press, 2022, pp. 99–158.
- 27 S. Tiwari, C. Singh and J. S. Singh, *Energy, Ecol. Environ.*, 2018, 3, 355-371.
- 28 J. S. Singh, A. Kumar, A. N. Rai and D. P. Singh, *Front. Microbiol.*, 2016, 7, 529.
- 29 D. R. Nayak, Y. J. Babu, A. Datta and T. K. Adhya, *J. Environ. Qual.*, 2007, **36**, 1577–1584.
- 30 R. Prasanna, V. Kumar, S. Kumar, A. K. Yadav, U. Tripathi, A. K. Singh, M. C. Jain, P. Gupta, P. K. Singh and N. Sethunathan, *Microbiol. Res.*, 2002, 157, 1–6.
- 31 L. Brennan and P. Owende, Renewable Sustainable Energy Rev., 2010, 14, 557–577.
- 32 Z. Chi, J. V. O'Fallon and S. Chen, *Trends Biotechnol.*, 2011, 29, 537–541.
- 33 M. G. De Morais and J. A. V. Costa, J. Biotechnol., 2007, 129, 439–445.
- 34 B. Wang, Y. Li, N. Wu and C. Q. Lan, *Appl. Microbiol. Biotechnol.*, 2008, **79**, 707–718.
- 35 L. Doron and C. A. Kerfeld, *Biochem. Soc. Trans.*, 2024, **52**, 997–1010.
- 36 A. Burlacot, O. Dao, P. Auroy, S. Cuiné, Y. Li-Beisson and G. Peltier, *Nature*, 2022, **605**, 366–371.
- 37 B. L. Montgomery, S. Lechno-Yossef and C. A. Kerfeld, *J. Exp. Bot.*, 2016, 67, 2931–2940.
- 38 L. Whitehead, B. M. Long, G. D. Price and M. R. Badger, *Plant Physiol.*, 2014, **165**, 398–411.
- 39 J. Wolfenden and T. A. Mansfield, *Proc. R. Soc. Edinburgh*, *Sect. B:Biol. Sci.*, 1990, **97**, 117–138.
- 40 I. Y. López-Pacheco, L. I. Rodas-Zuluaga, S. Fuentes-Tristan, C. Castillo-Zacarías, J. E. Sosa-Hernández, D. Barceló, H. M. Iqbal and R. Parra-Saldívar, J. CO<sub>2</sub> Util., 2021, 53, 101704.
- 41 M. Anjos, B. D. Fernandes, A. A. Vicente, J. A. Teixeira and G. Dragone, *Bioresour. Technol.*, 2013, **139**, 149–154.
- 42 V. Senatore, A. Buonerba, T. Zarra, G. Oliva, V. Belgiorno, J. Boguniewicz-Zablocka and V. Naddeo, *Chemosphere*, 2021, 273, 129682.
- 43 D. J. Farrelly, C. D. Everard, C. C. Fagan and K. P. McDonnell, *Renewable Sustainable Energy Rev.*, 2013, 21, 712–727.
- 44 B. Zhao and Y. Su, Algal Res., 2020, 51, 102066.
- 45 A. L. Gonçalves, Appl. Sci., 2021, 11, 871.
- 46 J. Bao, C. Zhuo, D. Zhang, Y. Li, F. Hu, H. Li, Z. Su, Y. Liang and H. He, *Plant Soil*, 2021, **463**, 97–112.
- 47 B. K. Chikkaswamy, Int. J. Adv. Res. Eng. Appl. Sci., 2015, 4, 1–15.

- 48 S. H. Marzouk, H. J. Tindwa, N. A. Amuri and J. M. Semoka, *Heliyon*, 2023, **9**, e13040.
- 49 D. Chittora, M. Meena, T. Barupal and P. Swapnil P, *Biochem. Biophys. Rep.*, 2020, 22, 100737.
- 50 J. U. Itelima, W. J. Bang, I. A. Onyimba, M. D. Sila and O. J. Egbere, *Direct Res. J. Agric. Food Sci.*, 2018, **6**, 73–83.
- 51 G. Berg, C. Zachow, H. Müller, J. Philipps and R. Tilcher, *Agronomy*, 2013, 3, 648–656.
- 52 B. Kour, P. Sharma, S. Ramya, S. Gawdiya, K. Sudheer and B. Ramakrishnan, *J. Appl. Phycol.*, 2024, **36**, 1859–1874.
- 53 International Energy Agency (IEA), World energy outlook 2023, Paris, 2023, https://www.iea.org.
- 54 B. B. Uzoejinwa, X. He, S. Wang, A. E. F. Abomohra, Y. Hu and Q. Wang, *Energy Convers. Manag.*, 2018, **163**, 468–492.
- 55 T. G. Ambaye, M. Vaccari, A. Bonilla-Petriciolet, S. Prasad, E. D. van Hullebusch and S. Rtimi, *J. Environ. Manage.*, 2021, 290, 112627.
- 56 D. Lee, H. Nam, M. W. Seo, S. H. Lee, D. Tokmurzin, S. Wang and Y. K. Park, *Chem. Eng. J.*, 2022, 447, 137501.
- 57 J. H. Hwang, J. Church, S. J. Lee, J. Park and W. H. Lee, *Environ. Eng. Sci.*, 2016, 33, 882–897.
- 58 R. A. Voloshin, M. V. Rodionova, S. K. Zharmukhamedov, T. N. Veziroglu and S. I. Allakhverdiev, *Int. J. Hydrogen Energy*, 2016, 41, 17257–17273.
- 59 D. Neupane, Bioengineering, 2022, 10, 29.
- 60 M. F. Demirbas, Appl. Energy, 2009, 86, S151-S161.
- 61 S. Joshi and S. Mishra, *Bioresour. Technol.*, 2022, 352, 127037.
- 62 S. J. Malode, K. K. Prabhu, R. J. Mascarenhas, N. P. Shetti and T. M. Aminabhavi, *Energy Convers. Manag.*, 2021, 10, 100070.
- 63 S. Hayashida, K. Ohta, P. Q. Flor, N. Nanri and I. Miyahara, Agric. Biol. Chem., 1982, 46, 1947–1950.
- 64 L. Qin, X. Zhao, W. C. Li, J. Q. Zhu, L. Liu, B. Z. Li and Y. J. Yuan, *Biotechnol. Biofuels*, 2018, 11, 1–10.
- 65 X. H. Hu, M. H. Wang, T. Tan, J. R. Li, H. Yang, L. Leach, R. M. Zhang and Z. Luo, *Genetics*, 2007, 175, 1479–1487.
- 66 R. A. Sheldon, ACS Sustain. Chem. Eng., 2018, 6, 32-48.
- 67 W. H. Chen, K. T. Lee and H. C. Ong, Energies, 2019, 12, 290.
- 68 A. Klanchui, N. Raethong, P. Prommeenate, W. Vongsangnak and A. Meechai, in *Network Biology*, Springer, Cham, 2017, pp. 75–102.
- 69 J. Dexter and P. Fu, Energy Environ. Sci., 2009, 2, 857-864.
- 70 M. Wang, G. Luan and X. Lu, J. Biotechnol., 2020, 317, 1-4.
- 71 P. Farrokh, M. Sheikhpour, A. Kasaeian, H. Asadi and R. Bavandi, *Biotechnol. Prog.*, 2019, **35**, e2835.
- 72 A. M. P. Anahas and G. Muralitharan, *Energy Convers. Manag.*, 2018, **157**, 423-437.
- 73 F. Andrews, M. Faulkner, H. S. Toogood and N. S. Scrutton, *Biotechnol. Biofuels*, 2021, **14**, 240.
- 74 L. Cepoi, N. Donţu, V. Şalaru and V. Şalaru, in Cyanobacteria for Bioremediation of Wastewaters, Springer, Cham, 2016, pp. 27–43.
- 75 M. Danouche, N. El Ghachtouli and H. El Arroussi, *Heliyon*, 2021, 7, e07609.
- 76 Y. Jia, W. Chen, Y. Zuo, L. Lin and L. Song, *Sci. Total Environ.*, 2018, **613**, 1324–1330.

77 T. Sun, H. Huo, Y. Zhang, Y. Xie, Y. Li, K. Pan, F. Zhang, J. Liu, Y. Tong, W. Zhang and L. Chen, *ACS Nano*, 2024, 18, 17694–17706.

**RSC Sustainability** 

- 78 H. I. El Shimi and S. S. Mostafa, *ARPN J. Eng. Appl. Sci.*, 2016, 11, 10259–10272.
- 79 A. Christodoulou and K. Stamatelatou, *Water Sci. Technol.*, 2016, 73, 453–462.
- 80 C. M. Monteiro, P. M. Castro and F. X. Malcata, *Biotechnol. Prog.*, 2012, **28**, 299–311.
- 81 D. Galinytė, G. Balčiūnaitė-Murzienė, J. Karosienė, D. Morudov, R. Naginienė, D. Baranauskienė, J. Šulinskienė, I. Kudlinskienė, A. Savickas and N. Savickienė, *Plants*, 2023, 12, 3150.
- 82 S. Shanab, A. Essa and E. Shalaby, *Plant Signal. Behav.*, 2012, 7, 392–399.
- 83 K. B. Chekroun, E. Sánchez and M. Baghour, *Int. J. Environ. Res. Publ. Health*, 2014, 1, 19–32.
- 84 Z. Zahra, Z. Habib, S. Hyun and M. Sajid, *Sustainability*, 2022, 14, 2041.
- 85 H. E. S. Touliabah, M. M. El-Sheekh, M. M. Ismail and H. El-Kassas, *Molecules*, 2022, 27, 1141.
- 86 I. Z. Ahmad, Lett. Appl. Microbiol., 2022, 75, 718-730.
- 87 A. C. Albertsson, C. Barenstedt and S. Karlsson, *Acta Polym.*, 1994, 45, 97–103.
- 88 P. Sarmah and J. Rout, *Environ. Sci. Pollut. Res. Int.*, 2018, 25, 33508-33520.
- 89 P. K. Roy, M. Hakkarainen, I. K. Varma and A. C. Albertsson, *Environ. Sci. Technol.*, 2011, 45, 4217–4227.
- 90 R. V. Kumar, G. R. Kanna and S. Elumalai, *J. Biorem. Biodegrad.*, 2017, 8, 2.
- 91 J. A. V. Costa, J. B. Moreira, B. F. Lucas, V. D. S. Braga, A. P. A. Cassuriaga and M. G. D. Morais, *Ind. Biotechnol.*, 2018, 14, 249–256.
- 92 M. Das and S. K. Maiti, J. Environ. Chem. Eng., 2021, 9, 105379.
- 93 S. Ansari and T. Fatma, PLoS One, 2016, 11, e0158168.
- 94 K. Meixner, I. Fritz, C. Daffert, K. Markl, W. Fuchs and B. Drosg, J. Biotechnol., 2016, 240, 61–67.
- 95 E. Rueda, E. Gonzalez-Flo, S. Mondal, K. Forchhammer, D. M. Arias, K. Ludwig, B. Drosg, I. Fritz, C. R. Gonzalez-Esquer, S. Pacheco and J. García, *Rev. Environ. Sci. Biotechnol.*, 2024, 23, 321–350.
- 96 T. Lopez-Arenas, M. González-Contreras, O. Anaya-Reza and M. Sales-Cruz, *Comput. Chem. Eng.*, 2017, **107**, 140–150.
- 97 D. Tiwari, N. Kumar, R. Bongirwar and P. Shukla, World J. Microbiol. Biotechnol., 2024, **40**, 263.
- 98 B. M. Berla, R. Saha, C. M. Immethun, C. D. Maranas, T. S. Moon and H. B. Pakrasi, *Front. Microbiol.*, 2013, 4, 246.
- 99 S. G. Hays and D. C. Ducat, *Photosynth. Res.*, 2015, **123**, 285–
- 100 M. Baunach, A. Guljamow, M. Miguel-Gordo and E. Dittmann, *Nat. Prod. Rep.*, 2024, 41, 347–369.
- 101 Y. Jeong, S. H. Cho, H. Lee, H. K. Choi, D. M. Kim, C. G. Lee, S. Cho and B. K. Cho, *Microorganisms*, 2020, 8, 1849.
- 102 J. Nogales, S. Gudmundsson and I. Thiele, *Bioengineered*, 2013, 4, 158–163.

- 103 G. Ma, H. Pei, W. Hu, X. Xu, C. Ma and X. Li, *Bioresour. Technol.*, 2014, **165**, 191–198.
- 104 D. Camsund and P. Lindblad, Front. Bioeng. Biotechnol., 2014, 2, 40.
- 105 T. Fujisawa, R. Narikawa, S. I. Maeda, S. Watanabe, Y. Kanesaki, K. Kobayashi, J. Nomata, M. Hanaoka, M. Watanabe, S. Ehira and E. Suzuki, *Nucleic Acids Res.*, 2017, 45, D551–D554.
- 106 Y. Kato, K. Inabe, R. Hidese, A. Kondo and T. Hasunuma, *Bioresour. Technol.*, 2022, 344, 126196.
- 107 P. Pachauri, P. Rai and S. Srivastava, in *Methods in Cyanobacterial Research*, CRC Press, 2024, pp. 187–204.
- 108 X. Li, C. R. Shen and J. C. Liao, Photosynth. Res., 2014, 120, 301–310.
- 109 J. Ungerer and H. B. Pakrasi, Sci. Rep., 2016, 6, 39681.
- 110 Y. N. Choi and J. M. Park, *Bioresour. Technol.*, 2016, **213**, 54–57
- 111 M. D. Deng and J. R. Coleman, *Appl. Environ. Microbiol.*, 1999, **65**, 523–528.
- 112 D. R. Nobles and R. M. Brown, Cellulose, 2008, 15, 691-701.
- 113 L. Liu, Y. Wang, N. Wang, X. Chen, B. Li, J. Shi and X. Li, *Biochem. Eng. J.*, 2021, **173**, 108070.
- 114 E. L. Lan and J. C. Liao, Metab. Eng., 2011, 13, 353-363.
- 115 R. Miao, X. Liu, E. Englund, P. Lindberg and P. Lindblad, *Metab. Eng. Commun.*, 2017, 5, 45–53.
- 116 X. Liu, H. Xie, S. Roussou and P. Lindblad, *Curr. Opin. Biotechnol.*, 2022, 73, 143–150.
- 117 H. Xie, J. Kjellström and P. Lindblad, *Biotechnol. Biofuels Bioprod.*, 2023, **16**, 134.
- 118 H. Luo, Y. Shi, F. Xie, T. Zhou, L. Gao, R. Yang and Z. Wang, *Ind. Crops Prod.*, 2023, **191**, 115976.
- 119 A. M. Varman, Y. Xiao, H. B. Pakrasi and Y. J. Tang, *Appl. Environ. Microbiol.*, 2013, **79**, 908–914.
- 120 P. Lindberg, S. Park and A. Melis, *Metab. Eng.*, 2010, **12**, 70–79.
- 121 J. Ungerer, L. Tao, M. Davis, M. Ghirardi, P. C. Maness and J. Yu, *Energy Environ. Sci.*, 2012, 5, 8998–9006.
- 122 C. Durall, P. Lindberg, J. Yu and P. Lindblad, *Biotechnol. Biofuels*, 2020, **13**, 1–13.
- 123 J. N. Markham, L. Tao, R. Davis, N. Voulis, L. T. Angenent, J. Ungerer and J. Yu, *Green Chem.*, 2016, 18, 6266–6281.
- 124 F. K. Bentley, A. Zurbriggen and A. Melis, *Mol. Plant*, 2014, 7, 71–86.
- 125 H. Kiyota, Y. Okuda, M. Ito, M. Y. Hirai and M. Ikeuchi, *Biotechnol. J.*, 2014, **185**, 1–7.
- 126 Y. Xue, Y. Zhang, S. Grace and Q. He, *J. Appl. Phycol.*, 2014, **26**, 219–226.
- 127 E. Żymańczyk-Duda, S. O. Samson, M. Brzezińska-Rodak and M. Klimek-Ochab, *Microorganisms*, 2022, **10**, 2318.
- 128 Y. Kato, K. Inabe, Y. Haraguchi, T. Shimizu, A. Kondo and T. Hasunuma, *Sci. Rep.*, 2023, 13, 7249.
- 129 S. Díaz-Troya, L. López-Maury, A. M. Sánchez-Riego, M. Roldán and F. J. Florencio, Mol. Plant, 2014, 7, 87–100.
- 130 A. Badary, S. Takamatsu, M. Nakajima, S. Ferri, P. Lindblad and K. Sode, *Mar. Biotechnol.*, 2018, **20**, 109–117.
- 131 S. Aikawa, A. Nishida, S. H. Ho, J. S. Chang, T. Hasunuma and A. Kondo, *Biotechnol. Biofuels*, 2014, 7, 1–8.

132 J. H. Jacobsen and N. U. Frigaard, *Metab. Eng.*, 2014, **21**, 60–70.

Critical Review

- 133 A. Joseph, S. Aikawa, K. Sasaki, Y. Tsuge, F. Matsuda, T. Tanaka and A. Kondo, *Biosci. Biotechnol. Biochem.*, 2013, 77, 966–970.
- 134 S. A. Angermayr and K. J. Hellingwerf, *J. Phys. Chem. B*, 2013, 117, 11169–11175.
- 135 S. A. Angermayr, A. G. Rovira and K. J. Hellingwerf, *Trends Biotechnol.*, 2015, 33, 352–361.
- 136 A. D. Van der Woude, S. A. Angermayr, V. P. Veetil, A. Osnato and K. J. Hellingwerf, *J. Biotechnol.*, 2014, **184**, 100–102.
- 137 S. Zhou, Y. Shao, N. Gao, L. Li, J. Deng, M. Zhu and S. Zhu, Sci. Total Environ., 2014, 482, 208–213.
- 138 I. S. Huang and P. V. Zimba, *Harmful Algae*, 2019, **86**, 139–209.
- 139 Y. Wang, T. Sun, X. Gao, M. Shi, L. Wu, L. Chen and W. Zhang, *Metab. Eng.*, 2016, 34, 60–70.
- 140 M. Hügler, C. Menendez, H. Schägger and G. Fuchs, *J. Bacteriol.*, 2002, **184**, 2404–2410.
- 141 S. Zhang, Y. Liu and D. A. Bryant, Metab. Eng., 2015, 32, 174–183.

- 142 R. Carpine and S. Sieber, *Curr. Res. Biotechnol.*, 2021, 3, 65–81.
- 143 T. Nawaz, L. Gu, S. Fahad, S. Saud, Z. Jiang, S. Hassan, M. T. Harrison, K. Liu, M. A. Khan, H. Liu and K. El-Kahtany, *Food Energy Secur.*, 2023, **12**, e495.
- 144 R. A. D. Al-Nedawe and Z. N. B. Yusof, *Microb. Bioact.*, 2023, **6**, 1–16.
- 145 A. Bouyahya, S. Bakrim, I. Chamkhi, D. Taha, N. El Omari, N. El Mneyiy, N. El Hachlafi, M. El-Shazly, A. Khalid, A. N. Abdalla and K. W. Goh, *Biomed. Pharmacother.*, 2024, 170, 115989.
- 146 A. Holland and S. Kinnear, Mar. Drugs, 2013, 11, 2239-2258.
- 147 J. P. Berry, M. Gantar, M. H. Perez, G. Berry and F. Noriega, *Mar. Drugs*, 2008, 6, 117–146.
- 148 G. Zanchett and E. C. Oliveira-Filho, *Toxins*, 2013, 5, 1896–1917.
- 149 R. P. Rastogi and A. Incharoensakdi, *Photochem. Photobiol. Sci.*, 2014, **13**, 1016–1024.
- 150 D. C. Ducat, J. A. Avelar-Rivas, J. C. Way and P. A. Silver, Appl. Environ. Microbiol., 2012, 78, 2660–2668.
- 151 N. A. Kamennaya, S. Ahn, H. Park, R. Bartal, K. A. Sasaki, H. Y. Holman and C. Jansson, in *Metabolic Engineering*, Elsevier, 2015, pp.76–85.