


Cite this: *RSC Adv.*, 2022, 12, 16396

Received 31st January 2022
Accepted 25th May 2022

DOI: 10.1039/d2ra00673a

rsc.li/rsc-advances

Biocatalytic conversion of sunlight and carbon dioxide to solar fuels and chemicals

Mandy Ching Man Yau,^a Martin Hayes^b and Shafeer Kalathil^{*a}

This review discusses the progress in the assembly of photosynthetic biohybrid systems using enzymes and microbes as the biocatalysts which are capable of utilising light to reduce carbon dioxide to solar fuels. We begin by outlining natural photosynthesis, an inspired biomachinery to develop artificial photosystems, and the rationale and motivation to advance and introduce biological substrates to create more novel, and efficient, photosystems. The case studies of various approaches to the development of CO₂-reducing microbial semi-artificial photosystems are also summarised, showcasing a variety of methods for hybrid microbial photosystems and their potential. Finally, approaches to investigate the relatively ambiguous electron transfer mechanisms in such photosystems are discussed through the presentation of spectroscopic techniques, eventually leading to what this will mean for the future of microbial hybrid photosystems.

Introduction

Initiated by the industrial revolution in the eighteenth century, our world has made major strides in several industries such as transport, research and development, and agriculture. However, coupled with the increase in human population, there is now a greater energy demand, consequently generating the current global warming crisis from rising carbon dioxide (CO₂)

emissions. Hydrogen (H₂) gas, a traditional energy carrier most commonly used in the production of fertilisers and refinement of oil, is a green alternative to reduce carbon emissions.^{1–3} It has a high energy density and a variety of applications but, along with carbon monoxide (CO), it is normally generated through the combustion of fossil fuels. Moreover, the biofuel is limited to H₂ *i.e.*, more complex biofuels such as ethanol and acetate cannot be made. Other current renewable technologies, such as solar and wind power, suffer from low efficiencies due to the fluctuations in the energy source input and this has created a large disparity between the supply and demand.^{4–6}

There are several pathways toward the synthesis of green, synthetic fuels to aid in the reduction of carbon emissions. This review aims to explore how biological hybrid systems can be

^aHub for Biotechnology in the Built Environment, Department of Applied Sciences, Faculty of Health and Life Sciences, Northumbria University, Newcastle, NE1 8ST, UK. E-mail: shafeer.kalathil@northumbria.ac.uk

^bJohnson Matthey Technology Centre, Cambridge Science Park, Milton Road, Cambridge, CB4 0FP, UK



Mandy Ching Man Yau obtained her MSc in Chemistry from Newcastle University. During her year abroad at Kyushu University, she conducted research on guest responsive luminescence of tetracyano Re(V) complexes under the supervision of Dr Masaaki Ohba. Working alongside Dr Fabio Cucinotta, she completed her thesis on tuning dye loadings in mesoporous silica. She is

currently undertaking a PhD at Northumbria University under the supervision of Dr Shafeer Kalathil on integrating light harvesting materials with microbes to generate biofuels from solar energy and waste carbon.



Martin Hayes is Global Technology Manager for the Life Science Technologies business at Johnson Matthey plc (JM). Martin has worked in a variety of R&D and customer facing technical roles across JM's catalyst businesses over the course of his career. He holds a PhD in heterogeneous catalysis from the University of Reading and recently graduated with an MBA from Durham University. His

interests lie in the application of catalytic technologies in sustainable processes for making the high value materials that enhance our quality of life. Martin was admitted as a Fellow of the Royal Society of Chemistry (FRSC) in 2017.



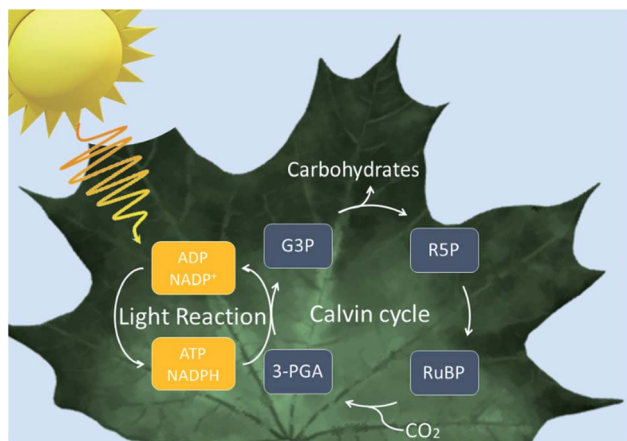


Fig. 1 Overview of the reactions within a natural photosynthetic system. Upon light absorption, electrons are excited twice by the photosystems to reduce nicotinamide adenine dinucleotide phosphate (NADP⁺) to NADPH. Adenosine triphosphate (ATP) is also generated upon water oxidation and electron transport which resulted in a proton gradient which drive ATP synthesis. In the dark reaction, CO₂ (and water) are combined with ribulose-1,5-biphosphate (RuBP) to yield 3-phosphoglycerate which is reduced to glyceraldehyde-3-phosphate with ATP and NADPH.

exploited to utilise CO₂ reduction to produce solar fuels and chemicals. While Reisner's article discusses similar topics in great detail,⁷ this updated review gives precedence to microbial hybrid photosystems and in particular, carbon dioxide fixation. This article also places the global importance of hydrogen as a fuel and touches upon the endeavours in the bioengineering, kinetics and proteomic and metabolomic aspect of microbial hybrid photosystems. Moreover, the potential for these microbial photosystems to be implemented on a commercial scale and how these biotechnologies are being supported is discussed.

Traditionally, artificial photosystems have high solar-to-chemical conversion efficiencies compared to natural photosynthetic systems, but they still have several cons – low stability,

high costs, and more importantly, they struggle to produce multicarbon products such as acetate.^{8,9} Microbial photohybrid system, on the other hand, is an emerging green technology and have great promise to overcome those challenges and become a viable alternative to current artificial photosystems.

To first understand how the idea of assimilating biocatalysts with inorganic semiconductors was first conceived, we must look at natural photosynthesis. Photoautotrophs, such as cyanobacteria and plants, utilise photosynthesis where solar energy and CO₂ are used to produce biomass. There are two phases to this reaction: light-dependent reaction and dark reaction. In the light-dependent reaction, light energy is used to make the reducing equivalents before proceeding to the dark reaction where they convert CO₂ into carbohydrates and other organic molecules *via* the Calvin cycle (Fig. 1).¹⁰ The overall efficiency for the conversion of solar energy to biomass is low due to energy losses occurring throughout the system.¹¹ However, the quantum efficiencies are high.^{12,13} Therefore, natural photosystems have been used as a blueprint for system design in current artificial and semi-artificial photosystems.

Artificial photosystems

Artificial photosystems have been commonly used to convert solar energy and feedstock chemicals such as CO₂ into fuels (Fig. 2). For example, Fujishima and Honda first demonstrated water splitting using solar energy and a photoelectrochemical (PEC) cell consisting of a titanium dioxide (TiO₂) electrode.¹⁴ Upon exposure to light, if the photon energy is equal to or greater than the bandgap of the material, electron-hole pairs are generated, and they can either recombine to generate heat or take part in photoreactions. Although PEC cells are theoretically able to achieve fairly high solar-to-chemical (STC) efficiencies, in the case of water oxidation, four electrons and four protons are lost to generate a O–O double bond ($498.36 \pm 0.17 \text{ kJ mol}^{-1}$).¹⁵ Thus, solar water splitting can be very energetically demanding with high kinetic and thermodynamic barriers. Nevertheless, to achieve high solar water-splitting efficiency, it is essential to develop a proton reduction catalyst capable of catalysing the production of H₂ *via* the conduction band edge potential of the semiconductor.

In comparison, CO₂ is more thermodynamically stable due to the presence of the strong C=O double bond ($532.2 \pm 0.4 \text{ kJ mol}^{-1}$).¹⁵ Hence, more amount of energy will be required to overcome the activation barrier. Similar to water splitting, upon light absorption, an electron-hole pair is generated and to favourably reduce CO₂, the choice of photocatalyst is extremely important. The photocatalyst should contain a suitable band structure where the conduction band edge must be more negative than the redox potential of CO₂ (-1.9 V vs. normal hydrogen electrode at pH 7) and the valence band edge more positive than water oxidation.^{16,17} If not, surplus photon energy will be lost as heat. Currently, there are limited materials with the suitable alignment of the energy band positions. Different strategies to alter the band gap have been employed including doping and surface modifications. Compared to plain bismuth oxide (Bi₂O₃), α -Bi₂O₃ doped with selenium had a smaller



Shafeer Kalathil received his PhD degree from Yeungnam University (with Prof. Moo Hwan Cho), and his doctoral research focused on electromicrobiology. He was a Japan Society for Promotion of Science (JSPS) Fellow at the University of Tokyo (with Prof. Kazuhito Hashimoto) and a Post-doctoral Fellow at KAUST (with Prof. Pascal Saikaly). He was a Marie Curie Individual Fellow at the University of Cambridge

(with Prof. Erwin Reisner) before moving to Northumbria University where he is currently a Vice Chancellor's Senior Fellow. His laboratory explores electrically active microbes to produce high-value chemicals and fuels from waste carbon.



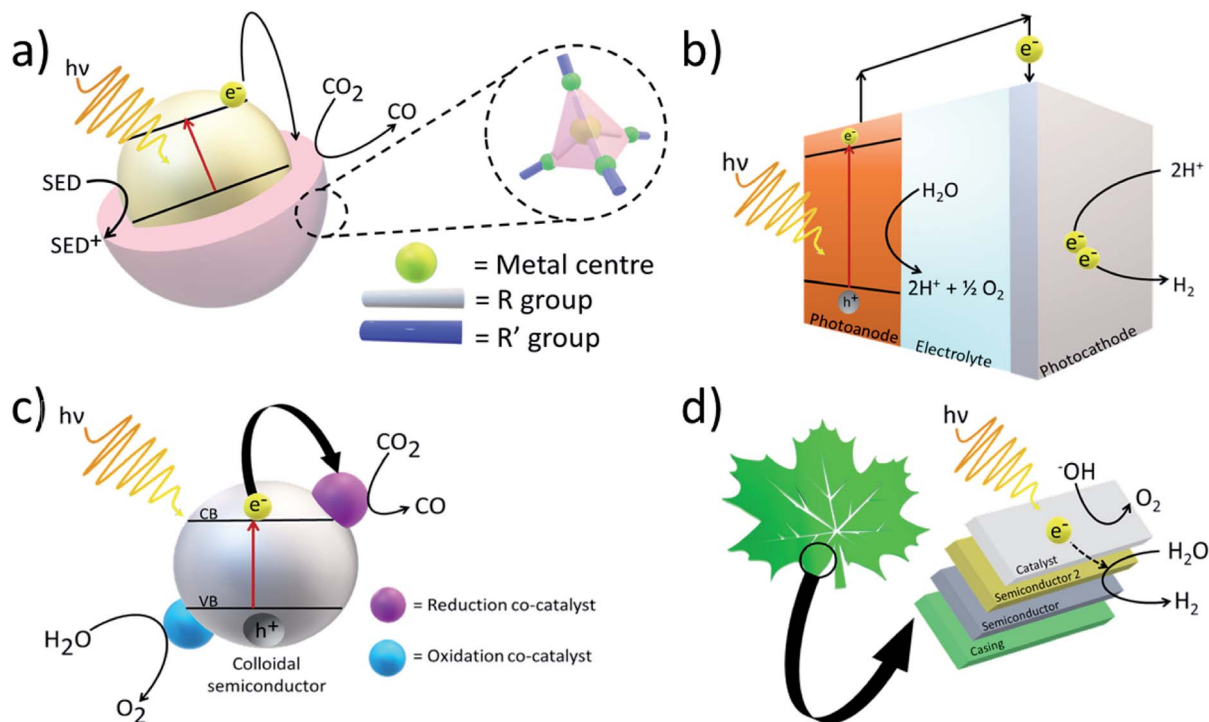


Fig. 2 Example artificial photosystems showing (a) a zeolitic imidazolate framework (ZIF) composed of tetrahedrally-coordinated transition metals and imidazolate linkers. ZIFs showing high adsorption and complexation towards CO₂ are combined with semiconductors with more appropriate bandgaps to generate electron-hole pairs for the photocatalytic reduction of CO₂. (b) a photoelectrochemical cell producing H₂ from solar energy (c) a general nanoparticle-based semiconductor for photocatalytic conversion of CO₂ to CO (d) artificial leaf composed of a tandem construct to produce H₂ from solar energy.

bandgap and increased photocatalytic activity.¹⁸ In a more novel approach, tuning the stoichiometric amount of formic acid added to a dicyandiamide polymerisation reaction led to the tailoring of the O- and N- link chains in the polymer which determined the band position shifts.¹⁹

The use of quantum dots (QDs) has also attracted lots of attention due to the quantum confinement effect giving rise to its distinct electronic properties, particularly its size-dependent bandgap.²⁰ H₂, CO, and methane (CH₄) were all able to be selectively retrieved in the photoreduction of CO₂ by tuning the size of the cadmium selenide (CdSe) crystals that had been doped onto nickel oxide.²¹ This in turn allowed for the control of the electron transfer kinetics. In another example, a zinc/cobalt-based ZIF was used to coat caesium lead bromide (CsPbBr₃) QDs to greatly enhance the photocatalytic reduction of CO₂ into CO and CH₄.²² Compared to bare CsPbBr₃, the electron consumption rate of CsPbBr₃-ZIF67 and CsPbBr₃-ZIF8 was 2.66 and 1.39 greater, respectively. *In situ* photoluminescence measurements on cobalt-containing ZIF revealed its role in promoting electron to cadmium sulfide (CdS).²³ In other studies, the potential of ZIFs in the activation of CO₂ has been demonstrated through enhanced activity.^{24–27}

Layered double hydroxides (LDH) have previously been investigated for the use of CO₂ reduction catalysts but the reported efficiencies have been low.^{28,29} By introducing palladium as a co-catalyst, a carbon nitride and LDH assembly proceeded to show an increase in reduction efficiency.³⁰ This was due to

the distortion of CO₂ upon interaction with the photocatalyst surface, thus allowing for high reactivity of the photocatalyst for CO₂ conversion. A similar improvement was also seen in the reduction of CO₂ to CH₄ through the use of sodium niobate (NaNbO₃) nanowires in which higher photocatalytic activity was observed compared to its bulk counterpart.³¹ However, the reason for the increase in photocatalytic activity in this system is most likely due to the surface-to-volume ratio and crystallinity.

Whilst artificial photosystems have greater solar to fuel efficiency compared to natural photosystems, there are still several drawbacks including high costs, toxicity, relatively low efficiencies, and inability to produce higher fuels – all of which are bottlenecks for commercialisation.^{6,32–34} Over the last few years, there has been a shift in focus towards the development of biological-based photosystems, *i.e.*, photobiohybrid systems. Their intrinsic metabolic pathways and numerous enzyme cascades enable the efficient conversion of simple feedstock into complex solar fuels and chemicals with greater product selectivity and stability. Photobiohybrid systems can be classified as enzymatic or microbial according to the biocatalyst.

Enzymatic photosystems

In a typical enzymatic photosystem, to which artificial photosystems are inspired, photocatalysts use solar energy to convert and generate photoexcited electrons which are then used to

convert reducing equivalents or are directly supplied to the enzyme (Fig. 3). The selective nature of enzymes limits the breadth of CO₂ transformations one can carry out. Therefore, although high rates and yields are achieved, sourcing the most suitable enzyme may result in a rise in costs. Redox enzymes play an important role in industrially relevant bioreactions and to carry out their jobs, reducing equivalents are generally required. Typically, nicotinamide adenine dinucleotide (NADH) and NADPH are generated by a secondary enzyme within the system. However, as mentioned above, photocatalysts can also provide reducing equivalents or directly transfer photoelectrons to active redox reactions. Photohybrid systems utilising enzymes have shown success in not only CO₂ reduction processes but also nitrogen-fixation and the synthesis of value-added products.³⁵

Hydrogenases, a type of metalloenzyme, are often used in the study of H₂ evolution due to their remarkable catalytic activity. In some cases, turnover frequencies of greater than 1000 s⁻¹ have been reported.^{36–38} [FeFe]-based hydrogenase was one of the first reported to be able to receive electrons from photocatalysts such as complexes involving rare metals and carbon nanotubes.^{39,40} Taking [FeFe]-hydrogenase from *Clostridium acetobutylicum*, Brown *et al.* integrated the enzyme with cadmium telluride (CdTe) nanocrystals but the random nature of absorption throughout the complex limited the efficiency of electron transfer complexes. This led to limited H₂ production.⁴¹

A common approach to producing formic acid from CO₂ is through the use of formate dehydrogenase (FDH) as a biocatalyst. Using FDH derived from *Saccharomyces cerevisiae* and zinc tetrakis(4-methylpyridyl)porphyrin as the light harvester, 62 μM formic acid was produced after 3 hours of irradiation.⁴² This was carried out in the presence of methyl viologen (MV) as the electron mediator. On the other hand, Noji *et al.* used tri(bipyridine)ruthenium(II) (Ru(bpy)₃²⁺) as the photosensitiser to produce formic acid from CO₂.⁴³ The use of a porous glass plate increased the conversion efficiencies of Ru(bpy)₃²⁺/MV²⁺/FDH from 1% to 22%. The nanopores within the porous glass plates allowed for the faster accumulation of MV^{•+} than in the solution, leading to greater interactions between FDH and MV^{•+} due

to the higher density. This is also previously seen when the rate of photoreduction was 16 times higher in a nanopore than in solution when photosystem II was employed as the photocatalyst.⁴⁴

In a hybrid system containing anatase/rutile TiO₂ mixture as the semiconductor, a ruthenium bipyridyl photosensitiser, and the enzyme carbon monoxide dehydrogenase, CO₂ was reduced to CO.⁴⁵ However, a SED was required to quench the hole left by the ruthenium complex and to regenerate the photosensitiser. Moreover, the average turnover rate with P25 TiO₂ was only 0.14 s⁻¹ at 20 °C which was lower than with a MV radical (100 s⁻¹). Even the addition of the electron donor ethylenediaminetetraacetic acid (EDTA) did not show any noticeable improvement, suggesting inefficient photosensitiser regeneration. A range of surface linkers was also tested to study the effects of enzyme binding to the TiO₂ surface with regard to electron transfer efficiency. However, those tested, including polymyxin B sulfate and glutamic acid (which had been previously described to strongly coordinate with TiO₂)⁴⁶ showed very little change in activity and stability. In one case, the addition of *o*-phosphorylethanol amine even resulted in a lower enzyme uptake and thus, decreased the rate of catalysis.

Microbial photosystems

Enzymatic inorganic hybrid systems appear to overcome the challenges presented by artificial photosystems. However, as described by the above studies, they suffer from several disadvantages including instability issues, poor durability, complicated experimental conditions, and high costs (Table 1). Out of the many carbon fixation pathways known, the Wood–Ljungdahl pathway is a classic for CO₂ fixation. In this approach, CO₂ is reduced to acetyl-coenzyme A and eventually converted to acetic acid which can be further used to make more complex products.^{47–51} By utilising the countless metabolic pathways present within the microbe and combining it with the high solar-to-energy efficiencies achieved by artificial semiconductors, products more complex and selective than what enzymes are capable of producing can be generated.



Fig. 3 Example of a typical enzymatic photohybrid system in which CO₂ is reduced by the enzyme to produce the appropriate biofuel.

Table 1 Summary of advantages and disadvantages of different photosystems

Type of photosystem	Advantages	Disadvantages
Non-biological	High solar energy conversion efficiencies Ease of design, tunability, and characterisation Broad absorption bands	Low product selectivity from CO ₂ reduction Tends to produce CO only pH gradients result in transfer limitations and may subject the electrodes to corrosion Internal resistance of the system may require an external bias to overcome Use of limited raw materials <i>e.g.</i> , rare earth metals
Enzymatic	High rates and yields High selectivity towards CO ₂ -to-chemicals	Expensive to isolate and purify Sensitive to a variety of environmental conditions (<i>e.g.</i> , O ₂ , pH and temperature) Prioritises survival over efficiency
Microbial	Utilises a myriad of enzyme cascades and metabolic pathways to synthesise complex products from simple feedstocks Resilient to environmental stress Self-healing capabilities Can be genetically engineered to increase conversion efficiencies, produce more complex chemicals, and improve product selectivity Mild reaction conditions	Susceptible to electron losses due to slow kinetics between microbe and electrode ROS produced during photocatalysis may deactivate microbial activities

In the following discussion, we summarise the recent achievements of artificial microbial photohybrid systems utilising solar energy and waste carbon to produce solar fuels and chemicals. Microbial photohybrid systems typically rely on the photosensitisation from organic or inorganic light absorbers to provide the reducing equivalents that allow them to carry out CO₂ reduction. Nanoparticles, in particular, are especially attractive and have drawn a lot of interest due to their quantum confinement effect. Essentially, if the size of the particle is comparable to that of the Bohr radius, then the optical and electronic properties become size-dependent.^{52,53} By altering the valence and conduction bands of the material, photocatalytic activity can be improved and in turn, greater solar-to-fuel efficiencies can be achieved.

Semiconductors are a vital aspect when striving towards an efficient photosystem capable of converting solar energy to biofuels. When discussing semiconductors, it is natural to immediately think about all those made in the laboratory. Decades of science and technology have made probing into light-harvesting capabilities of countless photosystems possible. However, long before we had the technology to investigate these systems, and before man-made semiconductors came about, nature had its inherent semiconductor found, remarkably, in rocks. In rock samples taken from China, Li *et al.* found that most of the samples had thin iron and manganese oxide coatings.⁵⁴ Presently, researchers often use Mn- and Fe- oxide in the study of photochemical systems due to their efficiency as photocatalysts and their intrinsic properties. As expected, it was found that photocurrent was only detectable in the Fe- and Mn- region on the coating and, upon exposure to solar energy, generated approximately 2.23×10^{16} photoelectrons. This is very meaningful as those photoelectrons could act as a major energy source to fuel the growth of photoelectronic

bacteria which has been reported numerous times.^{55,56} The study conducted by Li *et al.* is an important precursor to the research into artificial microbial systems as it reinforces the current research on the stimulation of bacterial growth obtained through phototrophic experiments using semiconductors as photocatalysts.⁵⁷

A pivotal development in the world of microbial photosystems was made by Sakimoto *et al.*, in the study of *Moorella thermoacetica* with CdS QDs.⁴⁷ From Tauc plots, a method of estimating bandgap energies, and absorption spectra, the quantum confinement effect was considered to have come into play when the direct bandgap was reported to be 2.51 ± 0.05 eV, as opposed to the expected lower value of 2.42 eV. The electrons generated by CdS when irradiated under visible light generated H⁺, a reducing equivalent, which was then used in the Wood–Ljungdahl pathway to produce acetic acid from CO₂ with a maximum yield of 90%. Later, TiO₂-manganese phthalocyanine (MnPc) was added to form a tandem system and cysteine/cystine (CySS/Cys) was chosen as the redox couple.⁵⁸ In general, this expanded hybrid system demonstrated greater acetic acid production. The system containing bare TiO₂ and *M. thermoacetica*-CdS had greater acetic acid production than *M. thermoacetica*-CdS but less than that of TiO₂-MnPc. These results demonstrate the superior efficiency of MnPc as a catalyst for CySS reduction. However, a recent study has highlighted pitfalls in the usage of organic chemicals such as cysteine as the SED.⁵⁹ The study revealed that bacteria can convert cysteine to acetate under dark condition which was largely neglected in previous studies due to poor control experiments. Hence, extreme care is needed in the selection of SED as bacteria can metabolize various substrates. Overall, this milestone, along with many other microbial photobiohybrid systems including the use of *Clostridium ljungdahlii* with CdS, *Escherichia coli* with Eosin Y,



Table 2 Example of hybrid photosystems

System	Biological substance	Light absorber	Reaction	Production rates/yields/activity	Comments	Reference
Inorganic	—	Magnesium–aluminium–LDH/carbon nitride	CO ₂ to CH ₄	3.7 μmol in 24 hours	—	30
	—	NaNbO ₃ nanowire	CO ₂ to CH ₄	653 ppm h ^{−1} g ^{−1}	—	31
	—	NaNbO ₃ bulk	CO ₂ to CO + H ₂	32 ppm h ^{−1} g ^{−1}	—	23
	—	Cobalt-ZIF-CdS	CO ₂ to CO + H ₂	124.4 μmol in 3 hours	TEOA = SED, bipyridine = e [−] transfer assistant, 82% CO selectivity	97% formate selectivity
Enzymatic	—	Lanthanum/rhodium-doped strontium titanium(III) oxide photosheet	CO ₂ to formate	0.08% solar to formate conversion efficiency	97% formate selectivity	45
	Carbon monoxide dehydrogenase I	Anatase/rutile TiO ₂ –RuP	CO ₂ to CO	0.14 s ^{−1}	2-(N-Morpholino) ethanesulfonic acid = SED	42
	Formate dehydrogenase	tetrakis(4-methylpyridyl) porphyrin	CO ₂ to HCOOH	62 μM in 3 hours	TEOA = SED MV ²⁺ = e [−] mediator	68
Microbial	<i>E. coli</i>	Silver indium sulfide/indium sulfide	H ⁺ to H ₂	1660 μmol in 3 hours	Cysteine = SED	67
	<i>M. thermoacetica</i>	Gold nanocluster	CO ₂ to acetate	6.01 mmol g ^{−1} over 7 days	Cysteine = SED	104
	<i>S. ovata</i>	Silicon nanowires	CO ₂ to acetate	0.3 g L ^{−1} over 7 days	—	69
	<i>R. palustris</i>	Graphitic carbon nitride	Fructose to PHB	6.73 g L ^{−1} after 96 hours	TEOA = SED	70
	<i>R. eutropha</i>	CdS nanorods	CO ₂ to PHB	28 mg over 48 hours	10% lactic acid = SED	73
		Graphitic carbon nitride catalyse	CO ₂ to PHB	41.02 mg L ^{−1} after 48 hours	—	71
		Core-shell QDs	PHB	100 mg g ^{−1} d ^{−1}	l-Ascorbic acid or HEPES = SED	80
	<i>R. capsulata</i>	Bi ₂ O ₃	2,3-Butanediol	10 mg g ^{−1} d ^{−1}	MV ²⁺ = e [−] mediator EDTA = SED	107
			Ethylene	1 mg g ^{−1} d ^{−1}		
<i>R. capsulata</i>	Bi ₂ O ₃	Isopropanol	2 mg g ^{−1} d ^{−1}	H ₂ = e [−] donor	64	
		H ⁺ to H ₂	~0.26 mL h ^{−1}			
<i>M. barkeri</i>	n ⁺ /p-Si/NiMo CdS	CO ₂ to CH ₄	17.6 mL over 72 hours	0.19 μmol h ^{−1} /13.70 μmol	Cysteine = SED	

and *Sporomusa ovata* with silicon nanowires have led to the innovation of many new and improved microbial hybrid photosystems (Table 2).

As can be inferred from the examples of microbial photo-biohybrid systems listed above, there are two approaches in which microbes can be interfaced with synthetic light absorbers: colloidal and non-colloidal. We begin the discussion below with colloidal microbial photosystems before moving on to non-colloidal systems which involve the use of nanowires and electrodes.

Microbial colloidal photosystems

In a typical colloidal photosystem, the semiconductor is present as a suspension. In some cases, they may be deposited onto the surface of the microbe. Such is the case for the microbial

photohybrid system consisting of CdS nanoparticles attached to the surface of *Clostridium autoethanogenum*.⁶⁰ Under light exposure, the CdS–*C. autoethanogenum* system produced almost 4-fold greater acetate than under dark conditions, and a 3-fold increase was observed under CO₂-only, confirming the H₂-driven autotrophic conversion to acetate. The system used ethanol to enable the solubility of lipoic acid in the culture medium. Later, it was confirmed that *C. autoethanogenum* oxidised ethanol due to the metabolic shift in response to the change in the redox environment. Ethanol oxidation enabled the hybrid system to obtain NADPH from the oxidised redox environment and this led to greater biomass production. Acetate production and NAD(P)H/NAD(P)⁺ increased when paired with H₂. NAD(P)H/NAD(P)⁺ play an important role in various biological processes, including biomass and biofuel production and often prevent efficient biotransformation



processes. Therefore, by enhancing the production of NAD(P)H/NAD(P)⁺, increased biomass generation will take place. However, microbial systems often contain various intricate biological networks and physiological systems and can self-organise their metabolic system, such that even with bioengineering, manipulation of the NAD(P)H/NAD(P)⁺ production pathway will be met with unexpected barriers.^{61–63} Future research into strategies to increase NADH/NAD⁺ or NADPH/NAD⁺ production will be required to understand and optimise conditions for NADH/NAD⁺ and NADPH/NAD⁺ production for various biotransformation processes. The CdS-*C. autoethanogenum* system demonstrates the viability of coupling microbes with semiconductors, and although further improvements are to be made in terms of electron capture for NADH and NADPH production, the system shows great potential for increased CO₂-fixing efficiency in nanoparticle-based hybrid photosystems through electron transfer processes.

The use of CdS nanoparticles has also been implemented in a hybrid system with *M. barkeri* which resulted in a CH₄ production rate of 0.19 μmol h⁻¹ with 0.34% quantum efficiency.⁶⁴ This system was further improved when doped with nickel, as well as CdS, which increased the photoelectron transfer within the system and led to an approximate rate of 0.24 μmol h⁻¹ CH₄.⁶⁵ Investigation into the proteomics of the system revealed that the addition of nickel caused a higher expression of proteins related to electron transfer, energy conversion and CO₂ fixation which contributed to the increased photoelectron transfer rate. Increasing protein expression is one method to increase electron transfer rates. Another viable method is through pi conjugation. Using the organic semiconductor poly(fluorene-co-phenylene) and perylene diimide derivative to photosensitise *M. thermoacetica*, the production of acetic acid from CO₂ was successful.⁶⁶ Efficient electron transfer was enabled through the pi conjugation present in the semiconductor which has been suggested to exhibit higher biocompatibility. Moreover, the quantum efficiency of the perylene diimide derivative was found to be comparable to inorganic semiconductors.

The production of acetic acid *via* CO₂ reduction was also realised through the use of gold nanoclusters with *M. thermoacetica*.⁶⁷ In a previous study carried out by Jiang *et al.*, the microbe viability before and after the photocatalytic reaction was measured through cell density and the lack of change indicated good viability of the cells.⁶⁸ In this study, the gold nanoclusters quenched reactive oxygen species (ROS) as well as acted as a light absorber. This enabled *M. thermoacetica* to ensure high bacterium viability over 6 days of continuous CO₂-fixation, showcasing the system's promising potential as a sustainable system.

Semi-artificial microbial systems can also be used to make more complex biofuels. Proof-of-concept was demonstrated in the works of Xu and coworkers where graphitic nitride and *R. eutropha* H16 were used to increase polyhydroxybutyrate (PHB) production from fructose 1.4-fold with triethanolamine (TEOA) as an SED.⁶⁹ Further works by the group include the use of CdS nanorods to produce PHB from CO₂ and fructose by *C. necator*.⁷⁰ Optimisation of CdS nanorods resulted in high photoactivity

and achieved a yield of 28 mg over 48 hours PHB from CO₂. On the other hand, Ding *et al.* employed different core-shell QDs to bind to specific enzyme binding sites in bacteria to improve product selectivity in photosynthetic biohybrid systems.⁷¹ Through this, not only was gram scale production of PHB achieved but also multi-carbon compounds. The production of PHB successfully increased using only a bioreactor with LED panels, illustrating the ease of scaling up beyond laboratory conditions and implementation on a commercial scale.

Many of the hybrid systems mentioned are limited by anaerobic conditions which may impede the assimilation of microbial hybrid photosystems in industry. The hybrid system presented by Liu *et al.* consists of CdS nanoparticles biologically precipitated onto the surface of *Thiobacillus thiooparus* and is able to reduce CO₂ to multicarbon glutamate synthase without being restricted by the need for an anaerobic environment.⁷² Interestingly, while chemically synthesised CdS predictably led to cell death in the later stages of cultivation, the biologically synthesised CdS showed a higher promoting effect for the growth of *T. thiooparus*. For the system to sustain cell growth in the presence of biologically synthesised CdS, not only is the electron transfer between the semiconductor-microbe interface efficient but also CO₂ fixation. Once CO₂ has been fixed by NADPH, the resulting precursors which undergo chemical transformations within *T. thiooparus* become biochemical components that support cell growth and replication. Hence, enabling the hybrid system to undergo self-replication. In not being limited by strict anaerobic conditions and its ability to sustain microbial growth with only simple, economical components, this work shows potential for the reduction of CO₂ in terms of design and sustainability.

While enzymes and microbes have individually been interfaced with light absorbers, the hybrid system established by Tremblay *et al.* involves both *R. eutropha* and H₂O₂-degrading catalyse for bioplastic production and water-splitting, respectively.⁷³ Although using the light absorber, graphitic nitride, and *R. eutropha* alone can successfully produce PHB, the yield was lower than with the addition of H₂O₂-degrading catalyse. Possible explanations for the increased yield include increased H₂ for *R. eutropha* to maintain its activity and the use of O₂ produced from water-splitting as an additional electron acceptor. The increased PHB production resulting from coupling water-splitting reactions and biochemical cascades within *R. eutropha* showcases a promising platform for efficient PHB production. However, compromises may have to be made with regards to performance and design due to the expense and temperament of enzymes.

H₂ is becoming an increasingly popular energy carrier, as evident by the global initiatives to invest in the hydrogen economy, such as the UK's Hydrogen Strategy,⁷⁴ USA's Hydrogen Program Plan,⁷⁵ Canada's Hydrogen Strategy,⁷⁶ Japan's Basic Hydrogen Strategy,⁷⁷ and the European Clean Hydrogen Alliance which consists of industries and authorities from various countries.⁷⁸ At present, the production of H₂ is still mainly from fossil fuels but in recent years, research into the electrolysis of water from renewable electricity has garnered a lot of attention. The downfall of this method, however, is that rare and precious



metals are used in the synthesis of the catalysts involved, meaning an increase in production costs and limited sustainability and scale-up potential.

As previously stated, current inorganic systems involving photovoltaic technology and semiconductors have surpassed natural biotransformation efficiencies. However, reaching similar conversion efficiencies is still a challenge yet to be achieved. *Clostridium butyricum* and *Escherichia coli*, along with semiconductors, were initially used to explore photocatalytic H₂ production.^{79–84} To this day, the pool of microorganisms used to explore H₂ production has expanded to include *Shewanella oneidensis* and more.^{85–89} In the works of Martins and coworkers, *Desulfovibrio desulfuricans*, *Citrobacter freundii*, and *S. oneidensis* were used in conjunction with CdS nanoparticles to investigate H₂ production.⁸⁴ *D. desulfuricans* was found to be the most efficient, owing to the high biological activity and/or efficient electron transfer. This was accompanied by the fact that it showed the highest hydrogenase activity compared to *S. oneidensis*, *E. coli*, and *C. freundii* (280 μmol g_{dcw}^{−1} min^{−1}, 8.4 μmol g_{dcw}^{−1} min^{−1}, 3.7 μmol g_{dcw}^{−1} min^{−1}, and 1.6 μmol g_{dcw}^{−1} min^{−1} respectively). High levels of the enzyme hydrogenase correlate with greater H₂ production as they catalyse H₂ formation from protons or the oxidation to protons. The high levels of hydrogenase in *D. desulfuricans* were found to be from the [NiFe] and [FeFe] families. These enzymes tend to be located in the periplasm of the microorganism,^{90,91} leading to a more efficient transfer of electrons from the nanoparticle to the enzyme than those found in the intracellular compartment. Cyanide was used to inhibit the hydrogenases within *D. desulfuricans*, almost completely preventing H₂ formation. This ultimately highlights the importance of hydrogenase in the pursuit of efficient H₂ formation. H₂ evolution was also possible through the use of a tandem involving silver indium sulfide/indium sulfide and *E. coli*.⁶⁸ In this system, the quantum efficiency of 3.3% was achieved and showed very little change in cell viability after the photocatalytic reaction, demonstrating the stability of the hybrid system.

A major bottleneck of microbial hybrid photosystems is the photoelectron transfer from the inorganic component of the system to the biological cell. In a study to increase H₂ production, reduced graphene oxide was chosen to be integrated with *S. oneidensis* MR-1.⁹² The reduced graphene oxide acted as a solid support surface and was able to efficiently collect electrons for the semiconductor, cuprous oxide (Cu₂O). In addition, the reduced graphene oxide operated as a pathway for electrons to travel between. All the above allowed for efficient electron transfer from the Cu₂O nanoparticles to *S. oneidensis* MR-1, thus contributing to efficient H₂ production, as evident in the increased photocatalytic H₂ production when compared to pristine Cu₂O and Cu₂O coupled with reduced graphene oxide (322.0 μmol g_{Cu₂O}^{−1} vs. 7.0 μmol g_{Cu₂O}^{−1} and 4.0 μmol g_{Cu₂O}^{−1} within 4 hours respectively).

It is important to elucidate the electron transfer pathway to understand the mechanism behind H₂ evolution within a hybrid system. Upon deletion of H₂ase, it was clear that within *S. oneidensis*, H₂ase was the active site for H₂ evolution as there was no H₂ production.⁹² Deletion of the proteins MtrC/OmcA

was found to have inhibited the production of H₂, thus establishing the fact that the cytochromes, including MtrA and MtrB, are involved in the electron transfer between the nanoparticle and *S. oneidensis*. While the work by Shen and coworkers concerns H₂ production, the understanding of the electron transfer pathway in the photocatalytic conversion extends to the production of all biofuels, not just H₂. In resolving and understanding the electron transfer pathway within a system, the overall design of hybrid systems can then be further improved, yielding greater electron transfer efficiency and ultimately, greater yields.

The kinetics of the electron transfer is extremely important in the conversion of solar energy to solar fuels. The previously mentioned work by Martin and coworkers on *D. sulfuricans* involved hydrogenases found in the periplasm of the microbe. Similarly, *S. oneidensis* also expresses periplasmic hydrogenases.⁹³ This was exploited by Shi and coworkers, in which copper indium sulfide/zinc sulfide (CuInS₂/ZnS) QDs were translocated into *S. oneidensis*, resulting in an increase in H₂ production due to the decrease in electron transfer distance and energy loss.⁹⁴ Through the deletion of certain genes, and the introduction of MV as a redox mediator, direct electron transfer from the QDs to *S. oneidensis* was inferred. As expected, single and double deletion of the periplasmic hydrogenase, [FeFe] and [NiFe], caused a decrease in the production of H₂, which suggests the two coenzymes play a crucial role in H₂ production. Deletion of the electron transport proteins had little to no effect on the production of H₂, nor did the introduction of MV as a redox mediator.

Manipulation of the genetic material within the microbe is becoming an increasingly popular method of increasing the production of biofuels. Using genetically engineered *E. coli* with precipitated CdS, [NiFe]-hydrogenase HyaABCDEF plasmid was transformed into the bacteria to catalyse H₂ formation.⁹⁵ Compared to non-HydA-induced hybrids, the resulting output per cell was much greater. In the case involving CdS and *E. coli*, genetic engineering was used to emphasise the expression of [FeFe]-hydrogenase gene, consequently yielding a photoactive *E. coli* hybrid system capable of converting solar energy into H₂ within a single cell.⁹⁶ Although the yield was less than previously reported studies involving TiO₂ and *E. coli*,⁹⁷ the genetically engineered hybrid is a viable method for enhancing photocatalytic H₂ production.

Microbial non-colloidal photosystems

Along with colloidal systems, microbes have also been interfaced with nanowires and PEC systems. Due to the large surface area, nanowires are extraordinary light harvesters. Initial studies on mammalian cells have demonstrated the ability to recognise and respond to nanoscale topographies.^{98–100} In response to this, a silicon nanowire array was interfaced with *S. oneidensis* MR-1.¹⁰¹ Through this model, it was discovered that not only attachment locations were influenced by the topography, but also the swimming patterns of the microbes. In a previous study conducted by Goto *et al.*, the tendency of single flagellated bacteria to travel in a straight trajectory was

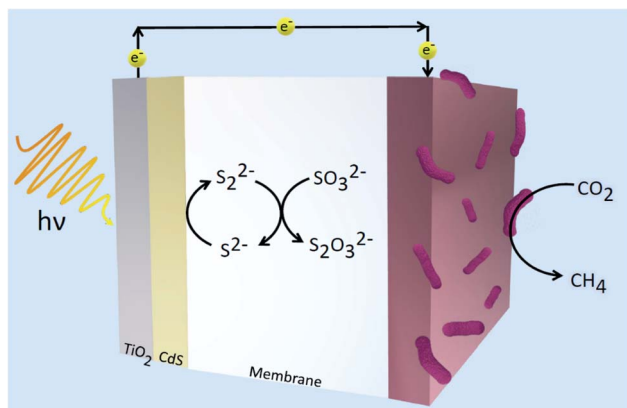


Fig. 4 Hybrid microbial photosystem designed by Xiao *et al.* consisting of a TiO₂/CdS photoanode and a biocathode for CO₂ reduction to CH₄.¹⁰⁶ The sulfide electrolyte solution was used as sacrificial reagents to consume holes and prevent the corrosion of CdS.

revealed.¹⁰² However, *S. oneidensis* was shown to have circled the nanowire despite being a singly flagellated bacterium. Moreover, the reversible bacterial attachment was demonstrated after some of the bacteria were found to separate from the Si array following a short period. The swimming behaviour exhibited by *S. oneidensis* is also observed by *E. coli* K-12 strains.¹⁰³ By highlighting the important role of topography and its interaction with bacteria, knowledge that will serve to aid in the design of future microbial-nanowire interfaces was realised.

As expected from the above study by Jeong *et al.*, the use of a nanowire array proved to be more advantageous than that of a single nanowire.¹⁰¹ The use of arrays has been widely implemented in several research studies. Liu *et al.* cultured *Sporomusa ovata* with a silicon nanowire array passivated by TiO₂ protection layer, leading to the production of acetic acid from CO₂ after the incubation period.⁵⁷ Using dual light-absorbers allow the system to fully absorb solar energy, thus increasing photovoltage for the system. This is demonstrated by the high

faradaic efficiency (90%). This system was taken further and extended by allowing genetically engineered *E. coli* to activate acetate into acetyl-CoA, ultimately partaking in the biosynthesis of various complex molecules such as cadinene, PHB, and *n*-butanol. The study is especially important as the system itself generated a local anaerobic environment at the bottom of the nanowire array, encouraging the reduction of CO₂ under aerobic (21% O₂) conditions. Su *et al.* later used highly doped p⁺ silicon nanowire arrays and platinum wire as the counter electrode to replace the TiO₂ photoanode previously used.¹⁰⁴ By adjusting the electrolyte pH and increasing the buffering capacity, the group successfully achieved the formation of closed-packed silicon nanowires. The resulting close-packed bio-hybrid system obtained a solar-to-acetate efficiency of 3.8% and a current density of $0.65 \pm 0.11 \text{ mA cm}^{-2}$, as opposed to the energy conversion of 0.38% for acetic acid production and 0.35 mA cm⁻² photocurrent by Liu and colleagues.⁵⁷

Using methanogens in a PEC cell setup, a one-step transformation from CO₂ to CH₄ with a faradaic efficiency of up to 96% with only solar energy as the only energy input was achieved.¹⁰⁵ Solar-to-fuel conversion efficiency was found to be at 0.1%, approximately half of that of natural photosynthesis. The greatest limitation in the system is expected to be from the bandgap of the TiO₂ photoanode employed. TiO₂ is often used in designing photosystems, but its downfall is that it can only absorb a small range of wavelengths of solar energy. Thus, in expanding the absorption range of materials used in photosystems, the efficiency will likely increase. A combination of TiO₂/CdS was also used as a photoanode in the reduction of CO₂ by methanogens.¹⁰⁶ Fig. 4 illustrates the system constructed by Xiao and coworkers which demonstrated 94.4% faradaic efficiency and solar-to-fuel conversion efficiency of 1.28%.¹⁰⁶ Cu₂-ZnSn₄ was later employed to sensitise the photoanode. Due to the lower charge separation resistance, the photogenerated electron-hole separation was much more favoured and this led to greater CH₄ production. Suspending microorganisms in solution is also a viable method to catalyse redox reactions fuelled by solar energy. By interfacing the archaea

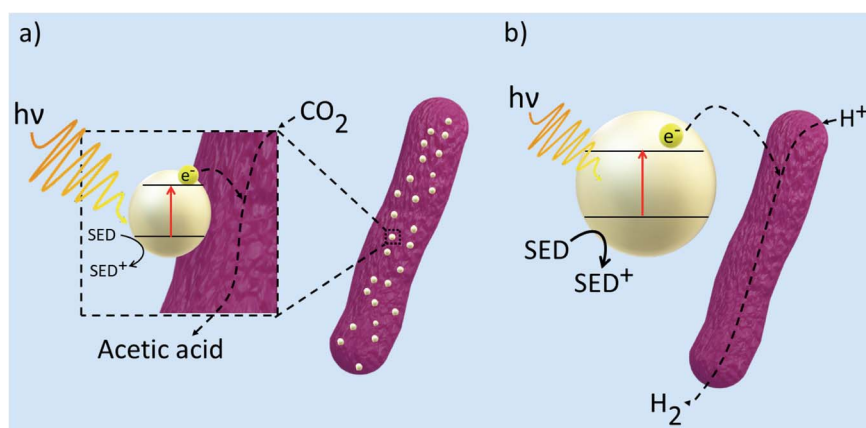


Fig. 5 General mechanism for electron transfer between a semiconductor and microbe in a hybrid photosystem. (a) Direct photoreduction of CO₂ to acetic acid in which the semiconductor is deposited onto the surface of the microbe (b) indirect photoreduction of CO₂ to H₂ by colloidal nanoparticles in the presence of electron mediators and SEDs.



Methanosarcina barkeri with a platinum electrode, Nichols *et al.* achieved a faradaic efficiency of up to 86% for electrochemical CO₂ to CH₄ conversion.¹⁰⁷ Replacing the platinum electrode with synthetic α -nickel sulfide H₂ evolution reaction catalyst resulted in similar faradaic efficiencies and CH₄ yields. Furthermore, by using an indium phosphide photocathode and a TiO₂ photoanode, the system proceeded to result in unassisted light-driven CH₄ production from CO₂ and this system yielded much greater CH₄ than that with the platinum electrode and α -NiS catalyst.¹⁰⁷ This proof-of-concept of bridging semiconductors with microbes proves efficient solar conversion to carbon-based fuels is attainable.

Elucidation of electron transfer mechanisms and microbial life in microbial photosystems

Although microbial hybrid systems have proven to be a promising approach to producing biofuels, achieving yields that can compete with current energy outputs from fossil fuels are still yet to be realised. Fig. 5 displays the general mechanisms taking place within a microbial hybrid photosystem but because of the knowledge gaps behind the mechanics of the hybrid system, the

finer details are still relatively unknown. Some challenges involve the seamless integration of synthetic and biological products and increasing the efficiencies of light absorption and solar-to-biofuel conversion. To work towards a system that can be applied to the industry, it is important to investigate the electron transfer mechanisms between the inorganic substrate and the biological component as they are the major bottleneck in the system. Therefore, at present, many techniques have been employed to probe into such interactions within the hybrid system to allow us to gain a deeper understanding.

Attenuated total reflection infrared (ATR-IR) (Fig. 6a) has been used to probe into the attachment of microbes onto inorganic surfaces in several previous works.^{108–111} ATR-IR spectroscopy is an attractive method for being not only suitable for solid samples but also for biological liquid samples. In the works of Freitag and colleagues, an acoustic trap was combined with ATR-IR spectroscopy to enable the entrapment of enzyme-labelled beads without the need for mechanical retention components, such as optical fibres, which are normally found in typical bead injection systems.¹¹² The set-up allowed for the conversion of *p*-nitrophenylphosphate into *p*-nitrophenol and phosphate under varying loadings of enzymes to be monitored, showcasing its great potential for enzymatic kinetic studies.



Fig. 6 (a) Schematics of ATR. The infrared beam interacts with the biofilm sample at one point of reflection (or at multiple points with multi-bounce ATR). Some of the evanescent waves formed from total internal reflection interact with the sample, resulting in an attenuated total reflection. Reactions can be monitored in real-time and information about kinetics and key transient reaction intermediates can be obtained, aiding in the study of kinetics and mechanism of the system. (b) Experimental setup of photoluminescence spectroscopy. The laser is directed to the sample using optical lenses. Once the laser hits the sample, the electrons are excited to higher energy states and as they relax, radiation is emitted which is detected by the photoluminescence spectrometer at specific wavelengths which reflect the energy level differences. The decay kinetics of the photoluminescent curve obtained will give information about the processes taking place in the hybrid photosystem. (c) In transient absorption spectroscopy, a pump pulse is used to excite the sample containing microbes interfaced with nanoparticles. Changes in the optical absorption of the sample are measured as a function of time and thus, the sample's absorbance is measured before and after pumping. The resulting information regarding the system's decay and lifetimes will shed light onto the kinetics of the system.

Steady-state photoluminescence (Fig. 6b) on the previously discussed work by Shen *et al.* was used to investigate the separation and transfer of photogenerated charge carriers in the system.⁹² In comparison with the absence of the Cu₂O nanoparticle, the steady-state photoluminescence intensity of Cu₂O/reduced graphene oxide decreased. This indicates that the separation of the photogenerated electron-hole pair was sufficient for a pathway between the Cu₂O nanoparticle and *S. oneidensis* cells for electron transfer. Similar observations were made with CuInS₂/ZnS QDs interfaced with genetically engineered *S. oneidensis*.⁹⁴

A more commonly used technique is time-resolved infrared spectroscopy (TRIR). TRIR is often used to study the kinetics, formation of intermediate products, and intramolecular charge transport of both biological and chemical molecular species.^{113–115} For example, TRIR spectroscopy, coupled with transient absorption (TA) spectroscopy (Fig. 6c), was used in an *M. thermoacetica*-CdS hybrid system to explain the decreasing CO₂ to acetic acid conversion rates with increasing H₂ incubation in the first three hours of photosynthesis.¹¹⁶ Changes in the vibrational modes of CO and carbon nitride, along with other amino acid residues, were observed and due to the peaks decaying on the same timescale as the TA signal, it was speculated the picosecond e[−] transfer was from a molecular process, not from a purely physical source.

In the same model, TA was used to aid in the elucidation of the decay kinetics of *M. thermoacetica*-CdS. It was reported that *M. thermoacetica*-CdS incubated with H₂ : CO₂ displayed faster decay kinetics compared to that incubated with glucose.^{47,116} The faster decay kinetics was attributed to the increased H₂ase activity and hence, the greater e[−] transfer rates and/or the greater quantity of e[−] acceptors in the H₂ incubated system. Unsurprisingly, multi-exponential decay in the timescale of picoseconds was found to be expressed in the hybrid system. Likewise, in the previously mentioned works of Luo *et al.*, when CuInS₂/ZnS QDs were integrated with *S. oneidensis* MR-1 through co-incubation, transient fluorescence and transient absorption lifetimes were shorter than that with CuInS₂/ZnS than with the hybrid system, indicating enzymes are efficient acceptors of photogenerated charge.⁹⁴ Decay kinetics in the same timescale as the *M. thermoacetica*-CdS hybrid was reported in a hybrid metal-semiconductor nanoparticle system consisting of CdSe-Au.¹¹⁷ However, lifetimes reported in the range of nanoseconds were found in Fe-Fe H₂ase-CdS systems,¹¹⁸ possibly owing to hydrogen bonding, solvent effects, and charge transfer barriers. From the evidence gathered *via* TA and TRIR spectroscopy, two proposed pathways for charge and energy transfer were proposed: the non-H₂ase-mediated pathway in glucose incubated cells and the membrane-bound H₂ase mediated pathways in H₂ incubated cells. Overall, the existence of a charge-transfer pathway was supported with TA and TRIR spectroscopy.

Due to the extremely fast nature of the charge separation and transfer, (from milliseconds to picoseconds), TA is also capable of providing information on parameters such as triplet excitons, exciton energy, and decay kinetics which can be used to speculate charge transfer processes.^{119–121} The previously mentioned

system consisting of cadmium selenium-gold (CdSe-Au) had undergone photoexcitation of the local surface plasmon resonance of the metal. By using ultrafast spectroscopy, along with high temporal resolution and near-infrared excitation, the plasmon-induced charge transfer (PICT) pathway was resolved, and it was discovered that hot-electron transfer to CdSe occurred in the timescale of fewer than 30 femtoseconds, whereas back transfer from the semiconductor to the metal was found to be within 210 fs.¹¹⁷ To compete with the extremely fast back-transfer to the metal, extraction of the electrons needs to be greater than the rate of back-transfer. This issue needs to be realised when designing hybrid systems to avoid inefficient and undesired transfer pathways. With suitable band alignments, such issues may be mitigated.

The kinetic study of this hybrid construct using high time resolution TA, along with the aforementioned studies, has highlighted the importance of spectroscopic techniques in the pursuit of elucidating electron transfer kinetics in photoactive systems. In establishing the requirements for resolving ultrafast photophysics of hybrid systems, there will be a greater understanding of the kinetic pathways within a system, resulting in a more intelligent design of hybrid systems which will allow for more diverse photocatalytic applications.

The involvement of proteomic and metabolomic analysis in the study of hybrid systems has recently become more prominent. Whilst the above-mentioned spectroscopic techniques will help clarify kinetics and electron transfer processes, proteomics and metabolomics will enable a greater understanding of the biocatalytic mechanisms, energy conservation in microbes, and physiological state of the organism. Taking the model *M. thermoacetica*-CdS hybrid system developed by Sakimoto and coworkers, Zhang *et al.* studied the energy conservation processes within the system using untargeted and targeted protein and metabolite quantification processes.¹²² Through this approach, it was discovered that the energy-metabolism-associated glycolysis and tricarboxylic acid cycle were active alongside the well-known Wood-Ljungdahl pathway. *M. thermoacetica* was also implied to contain an increased expression of proteins capable of protecting *M. thermoacetica* from oxidative stress but the damage far outweighed the protection provided.

Although the current biohybrid photosystems are still in their infancy, with the study presented by Zhang *et al.*,¹²² the aforementioned spectroscopic techniques, and more importantly, with technological advancements, the means to fabricate highly efficient, low cost, and industrially competitive biohybrid photosystems will become more tangible in the nearby future.

Conclusion

To conclude, while artificial photosystems and enzymatic photosystems have many advantages, artificial microbial photosystems present themselves as a much more powerful tool for converting sunlight and CO₂ to biofuels. However, there are still significant developments to be made. The commercialisation of hybrid photosystems continuously proves to be a challenge and several considerations including, but not limited to,



cost, toxicity, environmental impact, and the transition from laboratory to industrial scale still need to be addressed further.

With emerging technologies able to elucidate the electron transfer mechanisms in microbial hybrid photosystems, our understanding of the interactions occurring within the system will deepen, allowing for maximisation of solar-to-chemical yields, more complex solar fuel production, and smarter system design which will pave the way towards a revolutionised CO₂-utilisation technology sector. This will be further aided by the advancement of the synthetic biology sector, allowing for a new class of responsive, functionalised, and customisable hybrid systems to drive the future of green energy.

Future perspectives

The microbial conversion of CO₂ to fuels is promising but the variants of CO₂-fixation pathways all have different kinetic and ATP requirements. For example, the Calvin cycle is deemed inefficient due to its slow kinetics and high demand for ATP. For this reason, it is essential to design and develop pathways that can support greater carbon fixation. An example of this would be the pyruvate synthase-pyruvate carboxylate-glyoxylate pathway.¹²³ Genome-resolved metagenomics has also been used to identify potential enzymes that will show greater efficiency once integrated into CO₂-fixation pathways. Using metagenomics, Figueroa *et al.* identified genes responsible for phosphite oxidation and CO₂ reduction within *Candidatus Phosphitivorax anaerolimi* strain Phox-21.¹²⁴ It was discovered that while *Ca. P. anaerolimi* contained the genes necessary for Wood–Ljungdahl pathway, the genes for other carbon fixation pathways were absent. Moreover, CO₂ reduction *via* the Wood–Ljungdahl pathway was not feasible for Phox-21 as certain genes for particular carbonyl steps were missing. In light of these discoveries, Figueroa *et al.* proposed an approach as to how Phox-21 could reduce CO₂ to formate before proceeding to formate assimilation pathways. With the rapid development of analytical and biological techniques, carbon fixation may be enhanced to the standards of practical applications.

Expanding the library of microbes may also lead to unfamiliar reaction pathways that can be exploited to give greater STC efficiencies and more energy-dense solar fuels. The metabolic engineering of *Clostridium autoethanogenum* has already led to its large-scale industrial application for the production of ethanol from syngas.¹²⁵ Microorganisms lacking CO₂-fixing pathways, such as *E. coli* and *S. cerevisiae*, may be transplanted into heterotrophs like *S. oneidensis* and *G. sulfurreducens*. *Pyrococcus furiosus* is also another host candidate and attempts have been made to incorporate CO₂-fixation pathways into the archaea.^{126,127} The use of thermophile such as *P. furiosus* in hybrid microbial photosystems will be highly beneficial as solar transformations of CO₂ to volatile biofuels at higher temperatures will become more feasible.

Whilst the study and discovery of new carbon fixation pathways within the microbe is essential for more meaningful CO₂ transformation, investigations into the energy modules should not be disregarded. As described at the beginning of the review, a lot of energy is required to convert CO₂ to biofuels due

to its high stability. To circumvent this, the study of energy modules has been undertaken regarding semiartificial microbial photosystems. Owing to stoichiometric modelling, and computational analyses we now have a comparative analysis of autotrophic growth and production based on different natural and synthetic CO₂ fixation pathways and energy uptake.^{128,129} Advances in technology have now allowed for the multi-omics study of not only ATP production, but also electron transfer from semiconductor to microbe and CO₂ reduction which was carried out by Zhang *et al.*¹²²

Understanding the electron transfer between the semiconductor and microbe will also be essential for the future development of hybrid photosystems. Optimisation of the charge transfer processes through genetic engineering can take place, ensuring superior interaction between the biological and semiconductor interface. This in turn will theoretically give us greater STC efficiencies as well as a deeper understanding of how more complex biofuels can be generated.

Metal–organic frameworks (MOF) are becoming an emerging contender for light harvesting. MOFs are composed of transition metal ions that are connected to organic linkers and have crystalline, porous, extended structures. They are structurally tuneable, allowing for the assembly of higher surface areas for the capture of solar energy and its self-assembly into crystalline structures enables the MOF to be computationally analysed.^{130,131} MOFs have shown their capability through successfully harvesting water from desert air with only ambient sunlight as the energy input.^{132,133} For the solar conversion of CO₂ to biofuels, when interfaced with anaerobic microbes, MOFs have the potential to offer protection from oxygen and ROS, ultimately prolonging the lifetime of the bacteria and ultimately allowing for continuous biofuel production.¹³⁴ In using a zirconium nanocluster-based monolayer to wrap around *M. thermoacetica*, CO₂ transformation to acetate was able to last approximately 2.5 days whereas, without the MOF protection, CO₂ fixation was only able to last 1 day.¹³⁵ This was owed to the zirconium nanoclusters present within the MOF as it decomposed the hydrogen peroxide formed on the cell membrane, preventing ROS accumulation. Hydrogels have also been used to shield photosensitised *M. thermoacetica*, extending the viability of the bacterium and increasing acetate production.¹³⁶

Despite the benefits of microbial hybrid photosystems, commercialisation of large-scale production is still in the distant future. The major factors to consider are cost and efficiency, not to mention more research into novel artificial semiconductors that will bolster STC is needed. Hence, fossil fuels are still the most widely used source of fuel production.¹³⁷ However, initiatives such as the Paris Agreement and investments into green energy will help further the advancement of microbial biohybrid photosystems, especially as world leaders initiate the phasing out of fossil fuel industries. Like hybrid photosystems, next-generation materials consisting of smart, functional properties are being heavily invested in. In particular, synthetic biological materials are attracting attention for their potential in the construction of smart living due to their environmental responsiveness, adaptive abilities, self-healing



capabilities, and self-regeneration, just to name a few. Several successful engineering efforts have taken place to mimic these biological materials on a large scale. Buildings like the OME, an experimental biological house in the North-East of England which aims to test new, experimental biotechnologies for sustainable living¹³⁸ will help bridge the transition from laboratory to industry, and eventually, to real, everyday life. By monitoring the biological materials' response to the environment, the integration of microbial photosystems into these 'smart' properties will become more feasible, alleviating the population's dependency on fossil fuels.

In the 2000s, the invention of the genetic toggle switch first implemented within the plasmid of *E. coli* stimulated the synthetic biology sector and has now reached the fields of material science and engineering.¹³⁹ Rather than relying on bioengineering, the genetic toggle switch is controlled *via* manipulation of the network architecture. Demonstrating that the principles of engineering may be exploited to revolutionize simple cells into living machines, manipulation of the network architecture will enable the control of cell function and maximise STC efficiency. Moreover, by interfacing with inorganic materials such as gold nanoparticles and QDs, numerous environmentally responsive and controllable functional composite materials were developed. As previously mentioned, analysis of these environmental responses and adaption of the cellular functions will be important if the integration of microbial photosystems in buildings is to take place.

In recent years, the ability to transform waste carbon into biomass has become more and more important. Studies in this area include the works of Gleizer *et al.* who reported the engineering of *E. coli* to produce biomass from only CO₂.¹⁴⁰ To do this, the glycolysis and pentose-phosphate pathways, were disrupted whilst key enzymes in the Calvin cycle were expressed. A vital step in this study was the introduction of NAD-dependent FDH which would later be used to generate NADH to serve as the reducing power to drive carbon fixation. The strain was left to optimise for CO₂ fixation by allowing it to adapt and evolve for over 350 days. Genomic sequence analysis confirmed the presence of mutations that regulate gene expression and pathway regulations, overall achieving 100% biomass origin from CO₂ fixation. Similarly, Gassler and coworkers rewrote the xylulose monophosphate cycle to introduce the Calvin cycle.¹⁴¹ Methanol assimilating genes within the peroxisomes of *Pichia pastoris* were deleted and the energy and reducing power were supplied by methanol oxidation. These modifications resulted in the fungi utilising CO₂ as its sole carbon source, increasing its maximum growth rate on CO₂ more than two-fold. These studies highlight the importance of not only carbon fixation pathways but also the energy and reducing power on cell growth.

The works of Gleizer and co-workers have set an important milestone towards future works in the sustainable production of biofuels from CO₂ only, especially in allowing for a more in-depth study of how synthetic CO₂-fixation pathways can be assimilated into non-native hosts, as well as opening the likelihood of increasing the repertoire of microbes able to undergo solar reduction of CO₂ where new metabolic pathways may be

discovered. The accomplishments of Gleizer and more have stimulated the interdisciplinary field of synthetic biology and the various research currently in motion have demonstrated the great potential of smart materials in the application of sustainable production of biofuels in photohybrid systems.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by Research England (E3) Grant.

References

- 1 A. H. Awad and T. N. Veziroğlu, *Int. J. Hydrogen Energy*, 1984, **9**, 355–366.
- 2 M. Reese, C. Marquart, M. Malmali, K. Wagner, E. Buchanan, A. McCormick and E. L. Cussler, *Ind. Eng. Chem. Res.*, 2016, **55**, 3742–3750.
- 3 M. Balat, *Int. J. Hydrogen Energy*, 2008, **33**, 4013–4029.
- 4 P. Li, *IEEE Nanotechnol. Mag.*, 2008, **2**, 13–18.
- 5 H. Lund, *Energy*, 2007, **32**, 912–919.
- 6 D. Kim, K. K. Sakimoto, D. Hong and P. Yang, *Angew. Chem., Int. Ed.*, 2015, **54**, 3259–3266.
- 7 X. Fang, S. Kalathil and E. Reisner, *Chem. Soc. Rev.*, 2020, **49**, 4926–4952.
- 8 H. Lu, Z. Wang and L. Wang, *ACS ES&T Engg*, 2021, DOI: [10.1021/acsestengg.1c00336](https://doi.org/10.1021/acsestengg.1c00336).
- 9 C. Du, X. Wang, W. Chen, S. Feng, J. Wen and Y. A. Wu, *Mater. Today Adv.*, 2020, **6**, 100071.
- 10 J. Barber and B. Andersson, *Nature*, 1994, **370**, 31–34.
- 11 J. R. Bolton and D. O. Hall, *Photochem. Photobiol.*, 1991, **53**, 545–548.
- 12 A. Melis, *Plant Sci.*, 2009, **177**, 272–280.
- 13 J. U. Grobbelaar, *J. Appl. Phycol.*, 2009, **21**, 519–522.
- 14 A. Fujishima and K. Honda, *Nature*, 1972, **238**, 37–38.
- 15 Y.-R. Luo, *Comprehensive Handbook of Chemical Bond Energies*, CRC Press, Boca Raton, 1 edn, 2007.
- 16 A. A. Khan and M. Tahir, *J. CO₂ Util.*, 2019, **29**, 205–239.
- 17 X. Jiao, K. Zheng, Z. Hu, Y. Sun and Y. Xie, *ACS Cent. Sci.*, 2020, **6**, 653–660.
- 18 R. Sharma, M. Khanuja, S. N. Sharma and O. P. Sinha, *Int. J. Hydrogen Energy*, 2017, **42**, 20638–20648.
- 19 Y. Wang, F. Silveri, M. K. Bayazit, Q. Ruan, Y. Li, J. Xie, C. R. A. Catlow and J. Tang, *Adv. Energy Mater.*, 2018, **8**, 1801084.
- 20 S. Baskoutas and A. F. Terzis, *J. Appl. Phys.*, 2006, **99**, 013708.
- 21 H. Cho, W. D. Kim, K. Lee, S. Lee, G.-S. Kang, H.-I. Joh and D. C. Lee, *Appl. Surf. Sci.*, 2018, **429**, 2–8.
- 22 Z.-C. Kong, J.-F. Liao, Y.-J. Dong, Y.-F. Xu, H.-Y. Chen, D.-B. Kuang and C.-Y. Su, *ACS Energy Lett.*, 2018, **3**, 2656–2662.
- 23 S. Wang and X. Wang, *Appl. Catal., B*, 2015, **162**, 494–500.



- 24 Y. Guo, H. Yang, X. Zhou, K. Liu, C. Zhang, Z. Zhou, C. Wang and W. Lin, *J. Mater. Chem. A*, 2017, **5**, 24867–24873.
- 25 C. M. Miralda, E. E. Macias, M. Zhu, P. Ratnasamy and M. A. Carreon, *ACS Catal.*, 2012, **2**, 180–183.
- 26 S. Dou, J. Song, S. Xi, Y. Du, J. Wang, Z.-F. Huang, Z. J. Xu and X. Wang, *Angew. Chem., Int. Ed.*, 2019, **58**, 4041–4045.
- 27 T. Jose, Y. Hwang, D.-W. Kim, M.-I. Kim and D.-W. Park, *Catal. Today*, 2015, **245**, 61–67.
- 28 N. Ahmed, Y. Shibata, T. Taniguchi and Y. Izumi, *J. Catal.*, 2011, **279**, 123–135.
- 29 K. Teramura, S. Iguchi, Y. Mizuno, T. Shishido and T. Tanaka, *Angew. Chem., Int. Ed. Engl.*, 2012, **51**, 8008–8011.
- 30 J. Hong, W. Zhang, Y. Wang, T. Zhou and R. Xu, *ChemCatChem*, 2014, **6**, 2315–2321.
- 31 H. Shi, T. Wang, J. Chen, C. Zhu, J. Ye and Z. Zou, *Catal. Lett.*, 2011, **141**, 525–530.
- 32 M. A. Quader and S. Ahmed, in *Clean Energy for Sustainable Development*, ed. M. G. Rasul, A. k. Azad and S. C. Sharma, Academic Press, 2017, pp. 91–140, DOI: [10.1016/B978-0-12-805423-9.00004-1](https://doi.org/10.1016/B978-0-12-805423-9.00004-1).
- 33 M. Sandroni, R. Gueret, K. D. Wegner, P. Reiss, J. Fortage, D. Aldakov and M. N. Collomb, *Energy Environ. Sci.*, 2018, **11**, 1752–1761.
- 34 N. Cox, D. A. Pantazis, F. Neese and W. Lubitz, *Interface Focus*, 2015, **5**, 20150009.
- 35 S. Sultana, P. Chandra Sahoo, S. Martha and K. Parida, *RSC Adv.*, 2016, **6**, 44170–44194.
- 36 M. W. Ratzloff, M. B. Wilker, D. W. Mulder, C. E. Lubner, H. Hamby, K. A. Brown, G. Dukovic and P. W. King, *J. Am. Chem. Soc.*, 2017, **139**, 12879–12882.
- 37 H. Land, P. Ceccaldi, L. S. Mészáros, M. Lorenzi, H. J. Redman, M. Senger, S. T. Stripp and G. Berggren, *Chem. Sci.*, 2019, **10**, 9941–9948.
- 38 W. Lubitz, H. Ogata, O. Rüdiger and E. Reijerse, *Chem. Rev.*, 2014, **114**, 4081–4148.
- 39 W. Wang, T. Yu, Y. Zeng, J. Chen, G. Yang and Y. Li, *Photochem. Photobiol. Sci.*, 2014, **13**, 1590–1597.
- 40 D. Svedruzic Chang, T. McDonald, Y.-H. Kim, J. Blackburn, M. Heben and P. King, *Interaction of [FeFe]-hydrogenases with single-walled carbon nanotubes*, SPIE, 2007.
- 41 K. A. Brown, S. Dayal, X. Ai, G. Rumbles and P. W. King, *J. Am. Chem. Soc.*, 2010, **132**, 9672–9680.
- 42 R. Miyatani and Y. Amao, *Biotechnol. Lett.*, 2002, **24**, 1931–1934.
- 43 T. Noji, T. Jin, M. Nango, N. Kamiya and Y. Amao, *ACS Appl. Mater. Interfaces*, 2017, **9**, 3260–3265.
- 44 T. Noji, K. Kawakami, J.-R. Shen, T. Dewa, M. Nango, N. Kamiya, S. Itoh and T. Jin, *Langmuir*, 2016, **32**, 7796–7805.
- 45 T. W. Woolerton, S. Sheard, E. Pierce, S. W. Ragsdale and F. A. Armstrong, *Energy Environ. Sci.*, 2011, **4**, 2393–2399.
- 46 A. D. Roddick-Lanzilotta and A. J. McQuillan, *J. Colloid Interface Sci.*, 2000, **227**, 48–54.
- 47 K. K. Sakimoto, A. B. Wong and P. Yang, *Science*, 2016, **351**, 74–77.
- 48 K. K. Sakimoto, *Biochemist*, 2017, **39**, 30–33.
- 49 K. K. Sakimoto, N. Kornienko, S. Cestellos-Blanco, J. Lim, C. Liu and P. Yang, *J. Am. Chem. Soc.*, 2018, **140**, 1978–1985.
- 50 G. Borrel, P. S. Adam and S. Gribaldo, *Genome Biol. Evol.*, 2016, **8**, 1706–1711.
- 51 S. W. Ragsdale and E. Pierce, *Biochim. Biophys. Acta, Proteins Proteomics*, 2008, **1784**, 1873–1898.
- 52 M. A. El-Sayed, *Acc. Chem. Res.*, 2004, **37**, 326–333.
- 53 J. Jasieniak, M. Califano and S. E. Watkins, *ACS Nano*, 2011, **5**, 5888–5902.
- 54 A. Lu, Y. Li, H. Ding, X. Xu, Y. Li, G. Ren, J. Liang, Y. Liu, H. Hong, N. Chen, S. Chu, F. Liu, Y. Li, H. Wang, C. Ding, C. Wang, Y. Lai, J. Liu, J. Dick, K. Liu and M. F. Hochella, *Proc. Natl. Acad. Sci. U. S. A.*, 2019, **116**, 9741.
- 55 A. Lu, Y. Li, S. Jin, X. Wang, X.-L. Wu, C. Zeng, Y. Li, H. Ding, R. Hao, M. Lv, C. Wang, Y. Tang and H. Dong, *Nat. Commun.*, 2012, **3**, 768.
- 56 A. Lu, Y. Li, X. Wang, H. Ding, C. Zeng, X. Yang, R. Hao, C. Wang and M. Santosh, *Precambrian Res.*, 2013, **231**, 401–408.
- 57 C. Liu, J. J. Gallagher, K. K. Sakimoto, E. M. Nichols, C. J. Chang, M. C. Y. Chang and P. Yang, *Nano Lett.*, 2015, **15**, 3634–3639.
- 58 K. K. Sakimoto, S. J. Zhang and P. Yang, *Nano Lett.*, 2016, **16**, 5883–5887.
- 59 L. Göbbels, A. Poehlein, A. Dumnitch, R. Egelkamp, C. Kröger, J. Haerdter, T. Hackl, A. Feld, H. Weller, R. Daniel, W. R. Streit and M. C. Schoelmerich, *Sci. Rep.*, 2021, **11**, 2139.
- 60 S. Jin, Y. Jeon, S. Jeon Min, J. Shin, Y. Song, S. Kang, J. Bae, S. Cho, J.-K. Lee, R. Kim Dong and B.-K. Cho, *Proc. Natl. Acad. Sci. U. S. A.*, 2021, **118**, e2020552118.
- 61 A. Kondo, J. Ishii, K. Y. Hara, T. Hasunuma and F. Matsuda, *J. Biotechnol.*, 2013, **163**, 204–216.
- 62 S. Aslan, E. Noor and A. Bar-Even, *Biochem. J.*, 2017, **474**, 3935–3950.
- 63 S. Stephens, R. Mahadevan and D. G. Allen, *Front. Bioeng. Biotechnol.*, 2021, **8**, DOI: [10.3389/fbioe.2020.610723](https://doi.org/10.3389/fbioe.2020.610723).
- 64 J. Ye, J. Yu, Y. Zhang, M. Chen, X. Liu, S. Zhou and Z. He, *Appl. Catal., B*, 2019, **257**, 117916.
- 65 J. Ye, G. Ren, L. Kang, Y. Zhang, X. Liu, S. Zhou and Z. He, *iScience*, 2020, **23**, 101287.
- 66 P. Gai, W. Yu, H. Zhao, R. Qi, F. Li, L. Liu, F. Lv and S. Wang, *Angew. Chem., Int. Ed.*, 2020, **59**, 7224–7229.
- 67 H. Zhang, H. Liu, Z. Tian, D. Lu, Y. Yu, S. Cestellos-Blanco, K. K. Sakimoto and P. Yang, *Nat. Nanotechnol.*, 2018, **13**, 900–905.
- 68 Z. Jiang, B. Wang, J. C. Yu, J. Wang, T. An, H. Zhao, H. Li, S. Yuan and P. K. Wong, *Nano Energy*, 2018, **46**, 234–240.
- 69 M. Xu, P.-L. Tremblay, L. Jiang and T. Zhang, *Green Chem.*, 2019, **21**, 2392–2400.
- 70 M. Xu, P.-L. Tremblay, R. Ding, J. Xiao, J. Wang, Y. Kang and T. Zhang, *Sci. Total Environ.*, 2021, **753**, 142050.
- 71 Y. Ding, J. R. Bertram, C. Eckert, R. R. Bommarreddy, R. Patel, A. Conradie, S. Bryan and P. Nagpal, *J. Am. Chem. Soc.*, 2019, **141**, 10272–10282.
- 72 G. Liu, F. Gao, H. Zhang, L. Wang, C. Gao and Y. Xiong, *Nano Res.*, 2021, DOI: [10.1007/s12274-021-3883-0](https://doi.org/10.1007/s12274-021-3883-0).



- 73 P.-L. Tremblay, M. Xu, Y. Chen and T. Zhang, *iScience*, 2020, **23**, 100784.
- 74 UK hydrogen strategy, <https://www.gov.uk/government/publications/uk-hydrogen-strategy/uk-hydrogen-strategy-accessible-html-version>, accessed 28/03/22.
- 75 S. McQueen, J. Stanford, S. Satyapal, E. Miller, N. Stetson, D. Papageorgopoulos, N. Rustagi, V. Arjona, J. Adams, K. Randolph, D. Peterson, L. Hill, M. Koleva, T. Reinhardt, E. Frye, R. Schrecengost, A. Kokkinos, J. Litynski, R. Conrad, G. Soloveichik, D. Tew, S. Litzelman, J. Vetrano, R. Onuschak, A. Hahn, E. Hsieh and R. Costa, *Department of Energy Hydrogen Program Plan*, United States, 2020.
- 76 Canada's Hydrogen Strategy, <https://nrcan.gc.ca/climate-change-adapting-impacts-and-reducing-emissions/canadas-green-future/the-hydrogen-strategy/23080#shr-pg0>, accessed 28/03/22.
- 77 Basic Hydrogen Strategy, https://www.meti.go.jp/english/press/2017/1226_003.html, accessed 28/03/22.
- 78 European Clean Hydrogen Alliance Members, <https://ec.europa.eu/docsroom/documents/48994>, accessed 28/03/22.
- 79 A. A. N. Krasnovskii, V. V. Nikandrov and S. A. Nikiforova, *Dokl. Akad. Nauk SSSR*, 1985, **285**, 6.
- 80 P. Maruthamuthu, S. Muthu, K. Gurunathan, M. Ashokkumar and M. V. C. Sastri, *Int. J. Hydrogen Energy*, 1992, **17**, 863–866.
- 81 B. Wang, C. Zeng, K. H. Chu, D. Wu, H. Y. Yip, L. Ye and P. K. Wong, *Adv. Energy Mater.*, 2017, **7**, 1700611.
- 82 K. Xiao, T. H. Tsang, D. Sun, J. Liang, H. Zhao, Z. Jiang, B. Wang, J. C. Yu and P. K. Wong, *Adv. Energy Mater.*, 2021, **11**, 2100291.
- 83 Y. Honda, H. Hagiwara, S. Ida and T. Ishihara, *Angew. Chem., Int. Ed.*, 2016, **55**, 8045–8048.
- 84 M. Martins, C. Toste and I. A. C. Pereira, *Angew. Chem., Int. Ed.*, 2021, **60**, 9055–9062.
- 85 F. Grein, A. R. Ramos, S. S. Venceslau and I. A. C. Pereira, *Biochim. Biophys. Acta, Bioenerg.*, 2013, **1827**, 145–160.
- 86 P.-H. Hsieh, Y.-C. Lai, K.-Y. Chen and C.-H. Hung, *Int. J. Hydrogen Energy*, 2016, **41**, 21685–21691.
- 87 H. Koku, İ. Eroğlu, U. Gündüz, M. Yücel and L. Türker, *Int. J. Hydrogen Energy*, 2002, **27**, 1315–1329.
- 88 S. Mohanraj, K. Anbalagan, P. Rajaguru and V. Pugalenth, *Int. J. Hydrogen Energy*, 2016, **41**, 10639–10645.
- 89 A. Selvaggi, C. Tosi, U. Barberini, E. Franchi, F. Rodriguez and P. Pedroni, *J. Photochem. Photobiol., A*, 1999, **125**, 107–112.
- 90 J. W. Peters, G. J. Schut, E. S. Boyd, D. W. Mulder, E. M. Shepard, J. B. Broderick, P. W. King and M. W. W. Adams, *Biochim. Biophys. Acta, Mol. Cell Res.*, 2015, **1853**, 1350–1369.
- 91 Y. Nicolet, C. Piras, P. Legrand, C. E. Hatchikian and J. C. Fontecilla-Camps, *Structure*, 1999, **7**, 13–23.
- 92 H. Shen, Y.-Z. Wang, G. Liu, L. Li, R. Xia, B. Luo, J. Wang, D. Suo, W. Shi and Y.-C. Yong, *ACS Catal.*, 2020, **10**, 13290–13295.
- 93 G. Meshulam-Simon, S. Behrens, A. D. Choo and A. M. Spormann, *Appl. Environ. Microbiol.*, 2007, **73**, 1153–1165.
- 94 B. Luo, Y.-Z. Wang, D. Li, H. Shen, L.-X. Xu, Z. Fang, Z. Xia, J. Ren, W. Shi and Y.-C. Yong, *Adv. Energy Mater.*, 2021, **11**, 2100256.
- 95 W. Wei, P. Sun, Z. Li, K. Song, W. Su, B. Wang, Y. Liu and J. Zhao, *Sci. Adv.*, 2018, **4**, eaap9253.
- 96 Y. Honda, Y. Shinohara, M. Watanabe, T. Ishihara and H. Fujii, *ChemBioChem*, 2020, **21**, 3389–3397.
- 97 Y. Honda, M. Watanabe, H. Hagiwara, S. Ida and T. Ishihara, *Appl. Catal., B*, 2017, **210**, 400–406.
- 98 R. Yan, J.-H. Park, Y. Choi, C.-J. Heo, S.-M. Yang, L. P. Lee and P. Yang, *Nat. Nanotechnol.*, 2012, **7**, 191–196.
- 99 W. Kim, J. K. Ng, M. E. Kunitake, B. R. Conklin and P. Yang, *J. Am. Chem. Soc.*, 2007, **129**, 7228–7229.
- 100 K. Chung, J. A. DeQuach and K. L. Christman, *Nano LIFE*, 2010, **1**, 63–77.
- 101 H. E. Jeong, I. Kim, P. Karam, H.-J. Choi and P. Yang, *Nano Lett.*, 2013, **13**, 2864–2869.
- 102 T. Goto, K. Nakata, K. Baba, M. Nishimura and Y. Magariyama, *Biophys. J.*, 2005, **89**, 3771–3779.
- 103 Y. Kinoshita, T. Ishida, M. Yoshida, R. Ito, Y. V. Morimoto, K. Goto, R. M. Berry, T. Nishizaka and Y. Sowa, *Sci. Rep.*, 2020, **10**, 15887.
- 104 Y. Su, S. Cestellos-Blanco, J. M. Kim, Y.-x. Shen, Q. Kong, D. Lu, C. Liu, H. Zhang, Y. Cao and P. Yang, *Joule*, 2020, **4**, 800–811.
- 105 Q. Fu, S. Xiao, Z. Li, Y. Li, H. Kobayashi, J. Li, Y. Yang, Q. Liao, X. Zhu, X. He, D. Ye, L. Zhang and M. Zhong, *Nano Energy*, 2018, **53**, 232–239.
- 106 S. Xiao, Z. Li, Q. Fu, Y. Li, J. Li, L. Zhang, Q. Liao and X. Zhu, *Chem. Eng. J.*, 2020, **390**, 124530.
- 107 E. M. Nichols, J. J. Gallagher, C. Liu, Y. Su, J. Resasco, Y. Yu, Y. Sun, P. Yang, M. C. Y. Chang and C. J. Chang, *Proc. Natl. Acad. Sci. U. S. A.*, 2015, **112**, 11461.
- 108 S. J. Parikh, F. N. D. Mukome and X. Zhang, *Colloids Surf., B*, 2014, **119**, 38–46.
- 109 M. J. McWhirter, A. J. McQuillan and P. J. Bremer, *Colloids Surf., B*, 2002, **26**, 365–372.
- 110 S.-Y. Kang, P. J. Bremer, K.-W. Kim and A. J. McQuillan, *Langmuir*, 2006, **22**, 286–291.
- 111 M. J. McWhirter, P. J. Bremer, I. L. Lamont and A. J. McQuillan, *Langmuir*, 2003, **19**, 3575–3577.
- 112 S. Freitag, B. Baumgartner, S. Tauber, C. Gasser, S. Radcl, A. Schwaighofer and B. Lendl, *Anal. Chem.*, 2019, **91**, 7672–7678.
- 113 S. Maiti, G. C. Walker, B. R. Cowen, R. Pippenger, C. C. Moser, P. L. Dutton and R. M. Hochstrasser, *Proc. Natl. Acad. Sci. U. S. A.*, 1994, **91**, 10360.
- 114 K. Wilke and H. D. Breuer, *J. Photochem. Photobiol., A*, 1999, **121**, 49–53.
- 115 A. Mezzetti and W. Leibl, *Photosynth. Res.*, 2017, **131**, 121–144.
- 116 N. Kornienko, K. K. Sakimoto, D. M. Herlihy, S. C. Nguyen, A. P. Alivisatos, C. B. Harris, A. Schwartzberg and P. Yang, *Proc. Natl. Acad. Sci. U. S. A.*, 2016, **113**, 11750.



- 117 F. V. A. Camargo, Y. Ben-Shahar, T. Nagahara, Y. E. Panfil, M. Russo, U. Banin and G. Cerullo, *Nano Lett.*, 2021, **21**, 1461–1468.
- 118 M. B. Wilker, K. E. Shinopoulos, K. A. Brown, D. W. Mulder, P. W. King and G. Dukovic, *J. Am. Chem. Soc.*, 2014, **136**, 4316–4324.
- 119 H. Satzger and W. Zinth, *Chem. Phys.*, 2003, **295**, 287–295.
- 120 S. A. Haque, S. Handa, K. Peter, E. Palomares, M. Thelakkat and J. R. Durrant, *Angew. Chem., Int. Ed.*, 2005, **44**, 5740–5744.
- 121 B. S. Veldkamp, W.-S. Han, S. M. Dyar, S. W. Eaton, M. A. Ratner and M. R. Wasielewski, *Energy Environ. Sci.*, 2013, **6**, 1917–1928.
- 122 R. Zhang, Y. He, J. Yi, L. Zhang, C. Shen, S. Liu, L. Liu, B. Liu and L. Qiao, *Chem*, 2020, **6**, 234–249.
- 123 A. Bar-Even, E. Noor, N. E. Lewis and R. Milo, *Proc. Natl. Acad. Sci. U. S. A.*, 2010, **107**, 8889–8894.
- 124 I. A. Figueroa, T. P. Barnum, P. Y. Somasekhar, C. I. Carlström, A. L. Engbrektson and J. D. Coates, *Proc. Natl. Acad. Sci. U. S. A.*, 2018, **115**, E92–E101.
- 125 P. Dürre and B. J. Eikmanns, *Curr. Opin. Biotechnol.*, 2015, **35**, 63–72.
- 126 H. Atomi, T. Sato and T. Kanai, *Curr. Opin. Biotechnol.*, 2011, **22**, 618–626.
- 127 M. W. Keller, G. J. Schut, G. L. Lipscomb, A. L. Menon, I. J. Iwuchukwu, T. T. Leuko, M. P. Thorgersen, W. J. Nixon, A. S. Hawkins, R. M. Kelly and M. W. W. Adams, *Proc. Natl. Acad. Sci. U. S. A.*, 2013, **110**, 5840–5845.
- 128 A. G. Fast and E. T. Papoutsakis, *Curr. Opin. Chem. Eng.*, 2012, **1**, 380–395.
- 129 N. R. Boyle and J. A. Morgan, *Metab. Eng.*, 2011, **13**, 150–158.
- 130 V. Bernales, M. A. Ortuño, D. G. Truhlar, C. J. Cramer and L. Gagliardi, *ACS Cent. Sci.*, 2018, **4**, 5–19.
- 131 C. E. Wilmer, M. Leaf, C. Y. Lee, O. K. Farha, B. G. Hauser, J. T. Hupp and R. Q. Snurr, *Nat. Chem.*, 2012, **4**, 83–89.
- 132 N. Hanikel, M. S. Prévot, F. Fathieh, E. A. Kapustin, H. Lyu, H. Wang, N. J. Diercks, T. G. Glover and O. M. Yaghi, *ACS Cent. Sci.*, 2019, **5**, 1699–1706.
- 133 H. Kim, S. Yang, S. R. Rao, S. Narayanan, E. A. Kapustin, H. Furukawa, A. S. Umans, O. M. Yaghi and E. N. Wang, *Science*, 2017, **356**, 430–434.
- 134 Z. Ji, H. Zhang, H. Liu, O. M. Yaghi and P. Yang, *Proc. Natl. Acad. Sci. U. S. A.*, 2018, **115**, 10582.
- 135 Z. Ji, H. Zhang, H. Liu, O. M. Yaghi and P. Yang, *Proc. Natl. Acad. Sci. U. S. A.*, 2018, **115**, 10582–10587.
- 136 S. Cestellos-Blanco, H. Zhang and P. Yang, *Faraday Discuss.*, 2019, **215**, 54–65.
- 137 F. Perera, *Int. J. Environ. Res. Public Health*, 2017, **15**, 16.
- 138 *Hub for Biotechnology in the Built Environment*, <https://bbe.ac.uk/>, accessed 30/03/22.
- 139 T. S. Gardner, C. R. Cantor and J. J. Collins, *Nature*, 2000, **403**, 339–342.
- 140 S. Gleizer, R. Ben-Nissan, Y. M. Bar-On, N. Antonovsky, E. Noor, Y. Zohar, G. Jona, E. Krieger, M. Shamshoum, A. Bar-Even and R. Milo, *Cell*, 2019, **179**, 1255–1263.
- 141 T. Gassler, M. Sauer, B. Gasser, M. Egermeier, C. Troyer, T. Causon, S. Hann, D. Mattanovich and M. G. Steiger, *Nat. Biotechnol.*, 2020, **38**, 210–216.
- 142 Q. Wang, J. Warnan, S. Rodríguez-Jiménez, J. J. Leung, S. Kalathil, V. Andrei, K. Domen and E. Reisner, *Nat. Energy*, 2020, **5**, 703–710.

