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# A review of nanostructured non-titania photocatalysts and hole scavenging agents for CO<sub>2</sub> photoreduction processes

Jeannie Z. Y. Tan \* and M. Mercedes Maroto-Valer

The imperative for the development of sustainable energy technologies to alleviate the heavy reliance on fossil fuels as well as to mitigate the serious environmental issues associated with CO<sub>2</sub> emission has fostered the development of solar fuels through CO<sub>2</sub> photoreduction. The well-documented TiO<sub>2</sub> and modified TiO<sub>2</sub>-based photocatalysts have been shown to photoreduce CO<sub>2</sub> into hydrocarbons. Meanwhile, there is also an increasing interest in the utilisation of non-titania based materials, namely metal sulphides, oxides, oxynitrides and nitrides, for CO<sub>2</sub> photoreduction. Distinct from other published reviews, we discuss here recent progress made in designing metal sulphide, oxide, oxynitride and nitride photocatalysts for CO<sub>2</sub> photoreduction through morphological changes, aiming at providing a systematic summary of non-titania based materials for CO<sub>2</sub> photoreduction. Furthermore, the introduction of hole scavengers in order to maximise the CO<sub>2</sub> photoreduction efficiency is also reviewed.

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## 1. Introduction

Fossil fuels are currently unrivalled for energy generation, and our existing infrastructure is built to handle fossil fuels for transportation, heating and electricity.<sup>1</sup> Our heavy reliance on fossil fuels results in annual emissions of 32 Gt of CO<sub>2</sub>.<sup>2</sup> This is likely to increase to 36–43 Gt by 2035, subject to policies governing CO<sub>2</sub> emissions and energy use, even with increasing renewable energy sources.<sup>3</sup> To mitigate these environmental issues as well as alleviate our dependence on fossil fuels, harvesting the seemingly infinite solar energy and storing it in the form of chemical fuels hold significant promise to address current and future energy demands. Moreover, the chemical industry and a vast amount of chemical products rely heavily on using fossil fuel feedstock. This further motivates the development of sustainable processes to generate fuels and chemical feedstock from water and CO<sub>2</sub> using solar energy. Such a process is akin to photosynthesis in nature, and therefore, it is referred to as the artificial photosynthesis.

Photoelectrocatalytic reduction of CO<sub>2</sub> in aqueous suspensions using semiconducting powders was first proposed by Inoue *et al.* in 1979.<sup>4</sup> Later in 1987, the photocatalytic reduction of CO<sub>2</sub> to CH<sub>4</sub> in the presence of H<sub>2</sub>O was proposed by Thampi *et al.*<sup>5</sup> Since then, an increasing number of studies on the photo(electro)catalytic reduction of CO<sub>2</sub> have been conducted (Fig. 1). Among these studies, almost 50% focused on the materials employed as photocatalysts for conversion of CO<sub>2</sub> under UV and/or visible irradiation. The rest of the studies

concentrated mainly on modelling or process development. The use of TiO<sub>2</sub> as a photocatalyst for CO<sub>2</sub> reduction has been extensively studied and has been reviewed elsewhere.<sup>6–10</sup> However, the lack of systematic studies of non-TiO<sub>2</sub> semiconducting materials, namely metal sulphides, oxides, oxynitrides and nitrides, for CO<sub>2</sub> photoreduction (CO<sub>2</sub>PR) has inhibited the development of these photocatalysts compared to titania-based photocatalysts.

Although different photocatalysts (*i.e.*, titania and non-titania based semiconductors) have been proposed in the literature, the overall CO<sub>2</sub>PR conversion remains low especially under sunlight irradiation, making the CO<sub>2</sub>PR system not practical for commercialisation. To further increase the efficiency of CO<sub>2</sub>PR, the introduction of scavenging agents into the CO<sub>2</sub>PR system has been proposed. However, so far, the introduction of hole scavenging agents has not been systematically studied, though studies started in the last century. Therefore, the necessity to systematically scrutinise the recent development of non-TiO<sub>2</sub> photocatalysts and hole scavenging agents for CO<sub>2</sub>PR is of great demand.

There are enormous scientific and technical challenges involved in making even the simplest fuel, H<sub>2</sub>, and even more so for carbon-based fuels by means of CO<sub>2</sub> photoreduction. Similar to other photocatalytic processes, solar-driven photocatalytic conversion of CO<sub>2</sub> in the presence of H<sub>2</sub>O to hydrocarbon fuels uses semiconducting materials to harvest solar energy and provides active sites to allow the photocatalytic conversion process to occur. The basic steps of the photocatalytic process can be summarised as follows:

(1) generation of charge carriers (electron–hole pairs) by semiconducting materials upon absorption of photons with appropriate energy from the irradiation of light,

Research Centre for Carbon Solutions (RCCS), Heriot-Watt University, Edinburgh EH14 4AS, UK. E-mail: j.tan@hw.ac.uk



(2) separation of charge carriers and their transportation to the surface of the photocatalyst, and

(3) chemical redox reactions between the charge carriers and the reactants.

CO<sub>2</sub>PR with H<sub>2</sub>O into fuels is illustrated in Fig. 2. TiO<sub>2</sub> was the first material used for CO<sub>2</sub>PR,<sup>5</sup> and since then it has been widely used because of its abundance, availability, high chemical stability, low cost and non-toxicity.<sup>12</sup> Despite the great effort made in the CO<sub>2</sub>PR using TiO<sub>2</sub> and its derivative materials, the efficiency of the process remains low,<sup>7</sup> mainly attributed to the following factors:

(a) Rapid recombination of photogenerated electron-hole pairs;<sup>10</sup>

(b) Mild reducing power;

• The potential of the conduction band electrons is only slightly more negative than the multi-electron reduction potentials of CO<sub>2</sub>, thus providing a very small driving force, whereas the potential of the valence band holes is much more positive than the water oxidation potential.<sup>7</sup>

(c) Limited visible light absorption due to the wide bandgap (3.0–3.2 eV) of TiO<sub>2</sub>.<sup>13,14</sup>

Strategies including doping,<sup>15,16</sup> coupling with semiconductors,<sup>17–19</sup> dye sensitizing,<sup>20,21</sup> surface modification<sup>22,23</sup> etc. have been extensively used to improve TiO<sub>2</sub> photocatalysts and are summarised elsewhere.<sup>9,14,24,25</sup> However, the two most commonly used methods for extending the absorption range to visible light, namely sensitization or doping, do not fully address the optical issue of wide bandgap materials. Sensitizing agents (e.g., dyes or quantum dots) often degrade when exposed to UV light and photogenerate oxidizing holes in TiO<sub>2</sub>.<sup>7</sup> Dopant atoms, on the other hand, can become the centers of charge recombination. Moreover, the additional energy states associated with foreign atoms are highly localized, resulting in suppressed charge mobility.<sup>27</sup> Hence, while TiO<sub>2</sub> remains a benchmark photocatalyst, there is a lot of interest in developing other materials for CO<sub>2</sub>PR, such as carbon-based semiconductors (e.g., graphene-based composites,<sup>28,29</sup> carbon nanotube composites,<sup>30</sup> g-C<sub>3</sub>N<sub>4</sub> based composites<sup>31–33</sup> and

hybrid organic–inorganic materials<sup>34–37</sup>) and other inorganic transition or main group metal oxides, sulphides, oxynitrides, and nitrides. Since the use of carbon-based semiconductors for CO<sub>2</sub>PR has been reviewed elsewhere,<sup>30,34,38,39</sup> these photocatalysts are not be discussed herein.

Inorganic semiconductors, namely metal oxides, sulphides, oxynitrides and nitrides, are among the first semiconductors



*Prof. M. Mercedes Maroto-Valer (FRSE, FICHEM, FRSC, and FRSA) is the Assistant Deputy Principal (Research & Innovation) and Director of the Research Centre for Carbon Solutions (RCCS) at Heriot-Watt University. She leads a multidisciplinary team of over 50 researchers developing novel solutions to meet the worldwide demand for energy. Her team's expertise comprises energy*

*generation, conversion and industry, carbon capture, conversion, transport and storage, emission control, low carbon fuels, and low-carbon systems. She has over 450 publications, of which she edited 4 books, and 32% of her publications are among the top 10% most cited publications worldwide. Her research portfolio includes projects worth ~£35m, and she has been awarded a prestigious European Research Council (ERC) Advanced Award. She obtained a BSc with Honours (First Class) in Applied Chemistry in 1993 and then a PhD in 1997 at the University of Strathclyde (Scotland). Following a one-year postdoctoral fellowship at the Centre for Applied Energy Research (CAER) at the University of Kentucky in the US, she moved to Pennsylvania State University in the US, where she worked as a Research Fellow and from 2001 as an Assistant Professor and became the Program Coordinator for Sustainable Energy. She joined the University of Nottingham as a Reader in 2005, and within 3 years she was promoted to Professor in Energy Technologies. During her time at Nottingham she was the head of the Energy and Sustainability Research Division at the Faculty of Engineering. In 2012, she joined Heriot-Watt University as the first Robert Buchan Chair in Sustainable Energy Engineering and has served as the head of the Institute for Mechanical, Processing and Energy Engineering (School of Engineering and Physical Sciences) and the pan-University Energy Academy. She is a member of the Directorate of the Scottish Carbon Capture and Storage (SCCS). She holds leading positions in professional societies and editorial boards, and has received numerous international prizes and awards, including the 2018 Merit Award Society of Spanish Researchers in the United Kingdom (SRUK/CERU), 2013 Hong Kong University William Mong Distinguished Lecture, 2011 RSC Environment, Sustainability and Energy Division Early Career Award, 2009 Philip Leverhulme Prize, 2005 U.S. Department of Energy Award for Innovative Development, 1997 Ritchie Prize, 1996 Glenn Award—the Fuel Chemistry Division of the American Chemical Society and 1993 ICI Chemical & Polymers Group Andersonian Centenary Prize.*



*Dr Jeannie Z. Y. Tan received her BSc. (2010) from Universiti Sains Malaysia and her Msc. (2013) from Zhejiang University. She obtained her PhD (2017) from the University of Melbourne and after that joined the Research Centre for Carbon Solutions at Heriot-Watt University as a Research Associate. She is currently working on an EPSRC-funded project to develop innovative solutions for solar fuel conversion. She currently has 19 publications that have been cited over 200 times. She is also a reviewer for 2 journals (RSC Nanoscale Advances and Journal of Electronics & Telecommunication).*



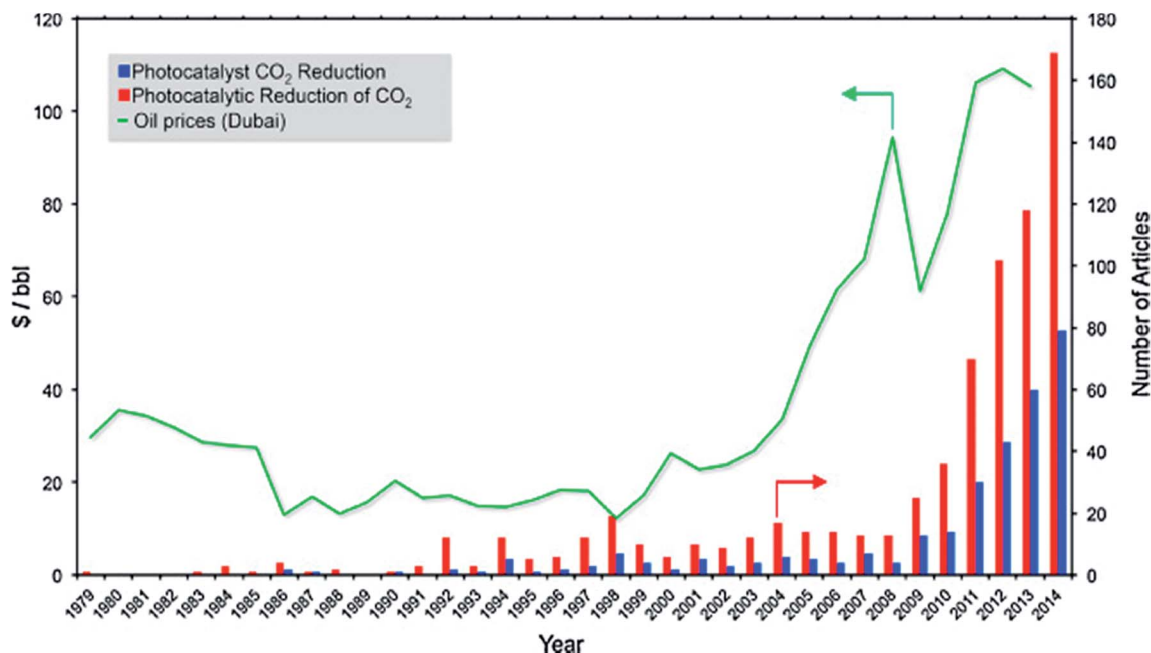


Fig. 1 Timeline for the number of articles published on CO<sub>2</sub> photoreduction vs. the price of oil. Reproduced from ref. 11 with permission.

used for solar-driven reactions. They possess relatively high stability, are low cost and absorb light consisting of photons with energy equal to or greater than their bandgap.<sup>40</sup> This very diverse group of materials includes both narrow and wide bandgap semiconductors; yet many of them offer a more favourable bandgap than TiO<sub>2</sub>. Moreover, many recent CO<sub>2</sub>PR developments follow similar trends to those for photocatalytic water splitting, as both processes share similar constraints on energy bands.<sup>41–43</sup> Specifically, the quest for new semiconductor materials is focused on the following points:<sup>27</sup>

- rising the valence band energy to decrease the bandgap,
- moving the conduction band to more reductive potentials,
- improving the quantum efficiency of exciton formation whilst suppressing charge recombination and

(d) using novel nanoscale morphologies to provide a large surface area with multiple photocatalytically active sites.

To achieve the quest mentioned above, different methods have been proposed previously and are reviewed in the following sections.

## 2. Non-TiO<sub>2</sub> materials for CO<sub>2</sub> photoreduction reactions

Although the position of conduction and valence bands is important for photocatalytic properties, the morphology of materials plays a critical role. Furthermore, manipulating the microstructure has also shown to alter the bandgap energy,<sup>44</sup> suppress the charge recombination,<sup>45</sup> enhance the diffusion of electrons towards the surface of photocatalysts,<sup>46</sup> induce quantum confinement effects<sup>47</sup> and provide more photocatalytic active sites, thereby enhancing the photocatalytic performance. In this section, nanostructured non-TiO<sub>2</sub> semiconducting materials for CO<sub>2</sub>PR published in the last two decades are reviewed, including metal sulphides, oxides, oxynitrides and nitrides.

### 2.1 Sulphides

Sulphide semiconductors received a lot of attention for CO<sub>2</sub>PR. This was because their valence band, made of 3p orbitals of the sulphur atoms, is located higher than those of their oxide analogues, resulting in the conduction band being more reductive.<sup>42</sup> Many sulphides have a narrow bandgap (*e.g.*, PbS and Bi<sub>2</sub>S<sub>3</sub>), with the absorption onset in the visible and infrared regions. Amongst sulphide semiconductors, ZnS and CdS were the most studied sulphides for CO<sub>2</sub>PR. ZnS is a wide bandgap semiconductor ( $E_g = 3.66$  eV in the bulk); however, it possesses

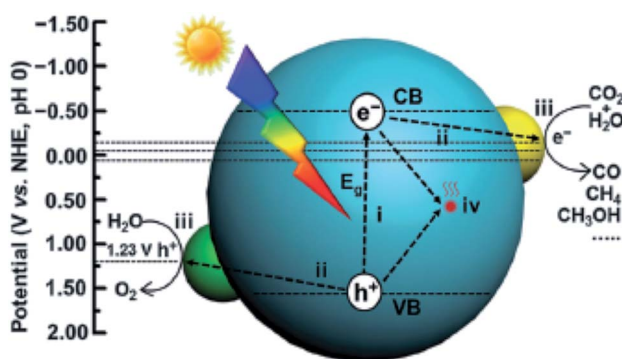


Fig. 2 Schematic illustration of photocatalytic CO<sub>2</sub> reduction with H<sub>2</sub>O over a heterogeneous photocatalyst. The dotted lines indicate the thermodynamic potentials for water oxidation and CO<sub>2</sub> reduction into CO, CH<sub>3</sub>OH and CH<sub>4</sub>. Reproduced from ref. 26 with permission.



a strong reducing power of the conduction band ( $E_{CB} = -1.85$  V vs. the NHE at pH 7).<sup>48</sup>

**Zinc-based materials.** The surface area of the photocatalyst is one of the key factors that can significantly affect the efficiency of the photocatalytic process. Kočí *et al.* proposed the immobilization of ZnS on montmorillonite, a representative natural clay mineral, which possesses a high surface area and layered structure, to optimise the efficiency of the CO<sub>2</sub>PR (Table 1 entry 1).<sup>49</sup> The study demonstrated that the amount of ZnS loaded affected the degree of agglomeration that consequently influenced the electronic configuration as well as the efficiency of the ZnS/montmorillonite nanocomposite in the CO<sub>2</sub>PR under UV irradiation (254 nm). A similar approach was demonstrated by Petra *et al.*, in which ZnS was loaded onto large-surface-area SiO<sub>2</sub> (340 m<sup>2</sup> g<sup>-1</sup>) to reduce CO<sub>2</sub> to formate using 2,5-dihydrofuran as the reducing agent.<sup>50</sup> The study revealed that the loading amount of ZnS significantly affected the yield and the optimal loading was 13% of ZnS into SiO<sub>2</sub>, resulting in 7 mmol g<sup>-1</sup> h<sup>-1</sup> of HCOOH. Nonetheless, the fabricated samples with coverages above 7% of ZnS on the SiO<sub>2</sub> matrix could suppress the photo-corrosion of ZnS to Zn(0), which is the major disadvantage of sulphides in an aqueous dispersion because the oxidation of lattice S<sup>2-</sup> ions leads to elemental sulphur and eventually to sulphate.<sup>51</sup>

Meng *et al.* proposed the co-doping of Cd and Cu into ZnS as one of the most active and optimised design routes for metal sulphide photocatalysts so far.<sup>52</sup> It was found that the doping of Cu could promote the formation of S vacancies and narrow the bandgap energy of ZnS, whereas surface modification of Cu-doped ZnS with Cd<sup>2+</sup> enhanced the product selectivity towards HCOOH (99%) under solar light irradiation. Recently, solid solutions of ZnLn<sub>2</sub>S<sub>4</sub> with a flower-like microstructure decorated with a cubic CeO<sub>2</sub> co-catalyst have been shown to exhibit enhanced CH<sub>3</sub>OH production (0.542 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup>) when compared to pristine CeO<sub>2</sub> and ZnLn<sub>2</sub>S<sub>4</sub> (0.139 and 0.073 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup>, respectively) under visible light irradiation ( $\lambda \geq 420$  nm).<sup>53</sup>

**Cadmium-based materials.** CdS (2.4 eV and the absorption onset at 520 nm) is a narrow bandgap metal sulphide photocatalyst. Hence, CdS suffers from rapid recombination of photogenerated electron-hole pairs. In order to enhance the separation of photogenerated electron-hole pairs, surface-phase junctions deduced by the same semiconductors were proposed. Chai *et al.* fabricated a mixed-phase CdS that is composed of wurtzite and zinc-blende crystalline phases recently (Table 1 entry 2).<sup>54</sup> The fabricated sample exhibited a long photogenerated electron lifetime and efficient charge transfer. The maximum CO and CH<sub>4</sub> evolution rate was 1.61 and 0.31 μmol h<sup>-1</sup> g<sup>-1</sup>, respectively, and these production rates were maintained even after 100 h.

The conduction band of CdS is less reductive ( $E_{CB} = -0.9$  V at pH 7 vs. NHE) than that of ZnS. Therefore, CdS is always decorated with noble metals, such as Ag. For instance, Zhu *et al.* proposed that the loaded Ag could act as an electron trap as well as an active site for CO<sub>2</sub>PR on CdS.<sup>55</sup> The photoproduction of CO was improved by three times when compared with that obtained with bare CdS. Alternatively, CdS can be supported with other

wide bandgap semiconductors to enhance its reducing power for CO<sub>2</sub>PR. Kisch *et al.* found that the coupling of CdS with ZnS strongly enhanced the CO<sub>2</sub>PR activity when compared to SiO<sub>2</sub>-supported CdS or ZnS samples because CdS and ZnS can absorb light at  $\leq 530$  nm and  $\leq 330$  nm, respectively.<sup>56</sup> The study reported that 5 wt% CdS loaded onto ZnS induced a 40-fold and 16-fold enhancement in the production of HCOOH ( $\sim 80$  mM,  $\lambda \geq 320$  nm, 3 h) when compared to unmodified CdS and ZnS, respectively. This strong enhancement was attributed to the electronic semiconductor-support interaction effect that improved the charge separation efficiency of the coupled semiconductor system. A similar observation was also reported by Kočí *et al.* recently, in which core-shell CdS/ZnS nanoparticles deposited on montmorillonite prepared by a one-pot synthesis exhibited enhanced CO<sub>2</sub>PR activity in water under UV irradiation ( $\lambda = 365$  nm).<sup>57</sup> The increase in the yield was due to the enhanced charge separation of CdS cores by ZnS shells, the increase of surface area and the inhibition of CdS photo-corrosion. CO<sub>2</sub>PR performed with CdS coupled with Bi<sub>2</sub>S<sub>3</sub>, having smaller bandgap energy than CdS, was also reported.<sup>59</sup> The Bi<sub>2</sub>S<sub>3</sub>/CdS nanocomposite fabricated with 15 wt% Bi<sub>2</sub>S<sub>3</sub> exhibited the highest methanol production from CO<sub>2</sub> (6.13 mmol g<sup>-1</sup> h<sup>-1</sup>, Table 1 entry 3), which was at least 50% higher than those obtained with bare Bi<sub>2</sub>S<sub>3</sub> (3.14 mmol g<sup>-1</sup> h<sup>-1</sup>) and CdS (2.01 mmol g<sup>-1</sup> h<sup>-1</sup>), under visible light irradiation. The enhanced photocatalytic activity suggested that the establishment of a heterojunction between CdS and Bi<sub>2</sub>S<sub>3</sub> could improve charge separation and subsequently prolong the lifetime of photogenerated electron-hole pairs. Moreover, the surface area of the Bi<sub>2</sub>S<sub>3</sub>/CdS nanocomposite, which was 24–27 m<sup>2</sup> g<sup>-1</sup>, was slightly higher than those of the bare CdS and Bi<sub>2</sub>S<sub>3</sub> (12 and 21 m<sup>2</sup> g<sup>-1</sup>, respectively). Hence, the synergistic effect of surface area and the heterojunction established between these two semiconductors had significantly improved the overall performance in CO<sub>2</sub>PR. Increasing the specific surface area does not only provide more active sites for the photocatalytic reaction, but also affects the optical properties of the material. For instance, Jin *et al.* recently proposed that by increasing the length-to-width ratio of Bi<sub>2</sub>S<sub>3</sub> nanoribbons, which increased the bandgap energy of Bi<sub>2</sub>S<sub>3</sub> from 1.22 to 1.38 eV, the CH<sub>3</sub>OH yield obtained was increased from 25.94 to 32.02 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup> under visible light irradiation ( $\lambda \geq 420$  nm).<sup>61</sup> However, the coupling of Bi<sub>2</sub>S<sub>3</sub> nanoribbons with CdS was not demonstrated. Hence, it will be interesting to see the performance of Bi<sub>2</sub>S<sub>3</sub> nanoribbons/CdS nanocomposites in the CO<sub>2</sub>PR.

The coupling of CdS with other metal oxides, such as WO<sub>3</sub>, has been demonstrated recently. For instance, Jin *et al.* proposed the coupling of WO<sub>3</sub> hollow spheres with CdS to form a hierarchical Z-scheme to increase the CO<sub>2</sub>PR efficiency.<sup>62</sup> The coupling of WO<sub>3</sub>-CdS had greatly enhanced the photo-conversion of CO<sub>2</sub> to CH<sub>4</sub> to  $\sim 1.0$  μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup> under visible irradiation ( $\lambda \geq 420$  nm), whereas pristine WO<sub>3</sub> and CdS only produced trace amounts of CH<sub>4</sub>.

Recently, the synthesis of Zn<sub>x</sub>Cd<sub>1-x</sub>S solid solutions has attracted extensive attention due to their versatility in tuning the band structures.<sup>63–65</sup> Moreover, the introduction of Zn can



Table 1 Photocatalytic CO<sub>2</sub> reduction yields obtained with various photocatalysts

| No.                | Photocatalyst   | Product(s) of CO <sub>2</sub> photoreduction<br>( $\mu\text{mol g}_{\text{catalyst}}^{-1} \text{h}^{-1}$ )                                | Light source  | Ref. |
|--------------------|---|---|---|------|
| <b>Sulphides</b>   |   |   |   |      |
| 1.                 | ZnS/montmorillonite nanocomposite   | CH <sub>4</sub> 1.17<br>CO 0.125  | UV 8 W Hg lamp ( $\lambda = 254 \text{ nm}$ )                 | 49   |
| 2.                 | CdS wurtzite/zinc-blende nanohybrid   | CO 1.61<br>CH <sub>4</sub> 0.31   | 300 W Xe lamp ( $\lambda \geq 420 \text{ nm}$ )               | 54   |
| 3.                 | Bi <sub>2</sub> S <sub>3</sub> /CdS   | CH <sub>3</sub> OH 6.13 $\text{mmol g}_{\text{catalyst}}^{-1} \text{h}^{-1}$  | 500 W Xe lamp ( $\lambda \geq 320 \text{ nm}$ )               | 59   |
|                    | CdS wurtzite/zinc-blende nanohybrid   | CO 1.61<br>CH <sub>4</sub> 0.31   | 300 W Xe lamp ( $\lambda \geq 420 \text{ nm}$ )               | 54   |
| 4.                 | Zn <sub>x</sub> Cd <sub>1-x</sub> S solid solution and tetra(4-carboxyphenyl)porphyrin iron(III) chloride                 | CO 1.28 $\mu\text{mol}$   | 300 W Xe lamp ( $420 \text{ nm} < \lambda < 780 \text{ nm}$ ) | 58   |
| 5.                 | Cu <sub>2</sub> S/CuS   | CH <sub>4</sub> $46.21 \pm 6.50 \mu\text{mol m}^{-2} \text{h}^{-1}$   | A.M 1.5 simulated sunlight                                    | 60   |
| 6.                 | RuO <sub>2</sub> -modified Cu <sub>x</sub> Ag <sub>y</sub> In <sub>z</sub> Zn <sub>k</sub> S <sub>m</sub> solid solutions | CH <sub>3</sub> OH 118.5  | 1000 W Xe lamp ( $\lambda > 400 \text{ nm}$ )                 | 69   |
| <b>Oxides</b>      |   |   |   |      |
| 7.                 | ZnO   | CH <sub>3</sub> OH 325  | 355 nm laser beam   | 74   |
| 8.                 | NiO   | CH <sub>3</sub> OH 388  |   |      |
| 9.                 | Fluffy mesoporous ZnO   | CO 0.73   | 8 W fluorescent tube ( $7 \text{ mW cm}^{-2}$ )               | 75   |
| 10.                | N-doped ZnO   | CO 0.04   |   |      |
| 11.                | ZnO plates  | CO 763.5 ppm $\text{g}_{\text{catalyst}}^{-1} \text{h}^{-1}$<br>CH <sub>4</sub> 205.2 ppm $\text{g}_{\text{catalyst}}^{-1} \text{h}^{-1}$ | 300 W Xe arc lamp   | 76   |
| 12.                | Ultralong and ultrathin single crystal Zn <sub>2</sub> GeO <sub>4</sub> nanoribbons                                       | CH <sub>4</sub> 25  | 300 W Xe arc lamp   | 77   |
| 13.                | Zn <sub>2</sub> GeO <sub>4</sub> nanorods   | CO 179 ppm $\text{g}_{\text{catalyst}}^{-1} \text{h}^{-1}$<br>CH <sub>4</sub> 35 ppm $\text{g}_{\text{catalyst}}^{-1} \text{h}^{-1}$      | 300 W Xe arc lamp   | 78   |
| 14.                | RuO <sub>2</sub> and Pt co-loaded Zn <sub>1.7</sub> GeN <sub>1.8</sub> O nano-sheaves                                     | ~55   | 300 W Xe arc lamp ( $\lambda > 420 \text{ nm}$ )              | 79   |
| 15.                | ZnGa <sub>2</sub> O nanosheet-scaffolded microspheres   | 69  | 300 W Xe arc lamp with an IR cut filter                       | 80   |
| 16.                | Zn <sub>2</sub> SnO <sub>4</sub> hexagonal nanoplates   | 47  | 300 W Xe arc lamp   | 81   |
| 17.                | Ce-doped ZnFe <sub>2</sub> O <sub>4</sub>   | CO ~20  | Visible light   | 82   |
| 18.                | Quasi-cubic WO <sub>3</sub>   | ~0.34   | 300 W Xe lamp   | 83   |
| 19.                | Ultrathin single crystal WO <sub>3</sub>  | ~1.1  | 300 W Xe arc lamp   | 84   |
| 20.                | Ultrathin W <sub>18</sub> O <sub>49</sub>   | CH <sub>4</sub> 2200  | Full-arc Xe lamp  | 85   |
| 21.                | Bi <sub>2</sub> WO <sub>6</sub> nanosheets with well-defined {001} facets   | 1.1   | 300 W Xe arc lamp   | 86   |
| 22.                | BiWO <sub>6</sub>   | CH <sub>3</sub> OH 32.6   | 300 W Xe lamp ( $\lambda > 420 \text{ nm}$ )                  | 87   |
| 23.                | NaNbO <sub>3</sub> nanowires  | CH <sub>4</sub> 653 ppm $\text{g}_{\text{catalyst}}^{-1} \text{h}^{-1}$   | 300 W Xe lamp   | 88   |
| 24.                | KNb <sub>3</sub> O <sub>8</sub> nanobelts   | CO 3.58   | 350 W Xe lamp   | 89   |
| 25.                | HNb <sub>3</sub> O <sub>8</sub> nanobelts   | CO 1.71   |   |      |
| 26.                | SrNb <sub>2</sub> O <sub>6</sub> nanorods   | CO 51.2   | 400 W Hg lamp   | 90   |
| 27.                | 3% NiO <sub>x</sub> -Ta <sub>2</sub> O <sub>5</sub> -1% immobilised on reduced graphene                                   | CH <sub>3</sub> OH 197.92   | 400 W metal halide lamp                                       | 91   |
| 28.                | Core-shell Ni/NiO-loaded N-InTaO <sub>4</sub>   | CH <sub>3</sub> OH 160  | Xe lamp (100 mW, $390 \leq \lambda \leq 770$ )                | 92   |
| 29.                | LaTa <sub>7</sub> O <sub>19</sub>   | CO 50   | 400 W Hg lamp   | 93   |
| 30.                | CaTa <sub>4</sub> O <sub>11</sub>   | CO 70   |   |      |
| 31.                | 1.0 wt% Ag-modified Ba-doped NaTaO <sub>3</sub>   | CO ~50  | 400 W Hg lamp   | 94   |
| 32.                | K <sub>2</sub> YTa <sub>5</sub> O <sub>15</sub>   | CO 91.9   | 4000 W Hg lamp  | 95   |
| 33.                | Ag-modified Ga <sub>2</sub> O <sub>3</sub>  | CO 10.5   | UV light  | 96   |
| 34.                | Lamellar BiVO <sub>4</sub>  | CH <sub>3</sub> OH 5.52   | 300 W Xe lamp (full spectrum)                                 | 97   |
| 35.                | CuGa <sub>1-x</sub> Fe <sub>x</sub> O   | CO ~9.2   | 300 W Xe arc lamp   | 98   |
| 36.                | CoAl-layered double hydroxides  | CH <sub>4</sub> 4.2   | 500 W Xe lamp   | 99   |
| <b>Oxynitrides</b> |   |   |   |      |
| 37.                | Porous TaON   | CH <sub>3</sub> CHO 0.52<br>C <sub>2</sub> H <sub>5</sub> OH 2.03   | 300 W Xe lamp   | 100  |
| 38.                | ZnAl <sub>2</sub> O <sub>4</sub> -modified ZnGa <sub>2</sub> ON   | CH <sub>4</sub> 9.2   | 300 W Xe lamp ( $\lambda \geq 420 \text{ nm}$ )               | 101  |
| <b>Nitrides</b>    |   |   |   |      |
| 39.                | GaN   | CO 1130<br>CH <sub>4</sub> 1.3  | 300 W Xe lamp   | 102  |
| 40.                | Rh/Cr <sub>2</sub> O <sub>3</sub> -decorated GaN nanowires  | CO 120<br>CH <sub>4</sub> 3.5   |   |      |



manipulate the structure of the surface atoms in CdS, which influences the adsorption or desorption of the reactants, intermediates and products in photocatalytic reactions.<sup>58</sup> In a very recent study, Li *et al.* integrated the well-defined floccule-like  $Zn_xCd_{1-x}S$  solid solution (Fig. 3) with tetra(4-carboxyphenyl)porphyrin iron(III) chloride for  $CO_2$  photoreduction under visible light irradiation.<sup>58</sup> The optimised photocatalyst, which was synthesized with  $Zn(NO_3)_2 \cdot 6H_2O$  and  $Cd(NO_3)_2 \cdot 4H_2O$  at 0.25 : 0.75 (ZCS-1, Fig. 3), produced 1.28  $\mu mol$  of CO with a selectivity of 93% after 4 h. However, pristine CdS and other synthesized  $Zn_xCd_{1-x}S$  solid solutions produced less than 0.4  $\mu mol$  of CO under visible light irradiation. The superior performance of ZCS-1 was attributed to the presence of sulphur vacancies that trapped photogenerated electrons, provided  $CO_2$  adsorption sites and facilitated the interaction between the  $Zn_xCd_{1-x}S$  solid solution and tetra(4-carboxyphenyl)porphyrin iron(III) chloride, resulting in efficient interfacial electron transfer for the subsequent photocatalytic reduction reaction.

**Copper-based materials.**  $Cu_{2-x}S$ , which have been shown to exhibit localised surface resonance in the near infrared region, and CuS, which has a direct bandgap of 2.0 eV, are nearly ideal for optimal sunlight absorption.<sup>66,67</sup> By carefully controlling the anodization voltage and temperature during the electrochemical anodization of copper foil and copper-coated Kapton substrates, in sodium sulphide electrolyte, copper sulphides with a nanowall nanostructure were obtained (Fig. 4).<sup>60</sup> The sample anodized with 1.5 V at 5 °C exhibited the highest methane formation in the  $CO_2$ PR (Table 1 entry 5) under the irradiation of simulated sunlight. At low voltage and temperature, sulphur diffusion was low, leading to a lower concentration of excess sulphur in the sample that yielded  $Cu_2S$ . As a result, less bulk Cu vacancy defects were formed within the sample.  $Cu_2S$  exhibited higher charge mobility than the CuS nanostructured array, which was obtained at high temperature and high voltage.

To engineer the bandgap energy of the photocatalyst that matches the solar spectrum, a solid solution with large and small bandgap semiconductors was proposed. For instance, Arai *et al.* used the Cu-based sulphide complex  $Cu_2ZnSnS_4$  with a direct bandgap of 1.5 eV and a large optical absorption coefficient and obtained a high selectivity of the photoelectrochemical  $CO_2$  reduction reaction (>80%).<sup>68</sup> The Cu-based sulphide complex reported by Liu *et al.* showcased that the Cu-based sulphide complex was able to reduce  $CO_2$  under visible light irradiation in the presence of a Ru co-catalyst.<sup>69</sup> The  $Ru-Cu_xAg_yIn_zZn_kS_m$  solid solutions induced the formation of methanol in  $CO_2$ PR under visible light irradiation (Table 1 entry 6). Although the study reported that the optimal performance could be obtained through the elemental composition manipulation, the nanostructures of the sulphide complex were not revealed. It is therefore questionable whether the efficiency of these photocatalysts could be further enhanced through the manipulation of their microstructures. Moreover, the stability of metal sulphates in most of the studies has not been demonstrated, and this should be emphasized more in future work.

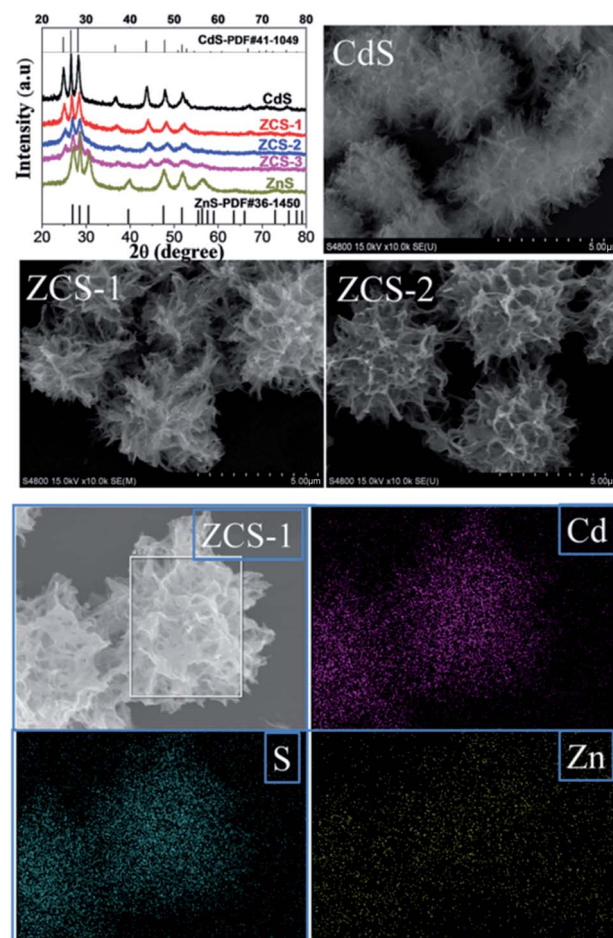


Fig. 3 XRD patterns, SEM images and EDX mapping of the synthesized CdS and  $Zn_xCd_{1-x}S$  solid solutions. Reproduced from ref. 58 with permission.

## 2.2 Oxides

Semiconducting oxides have been widely used as photocatalysts because of their stability and resistance to photocorrosion under irradiation. Hence, oxides have been used for photooxidation and photoreduction reactions. The intrinsic properties of metal oxides play a critical role in determining their feasibility for  $CO_2$ PR. For example,  $CO_2$ PR was observed for p-type NiO covalently linked with a Zn porphyrin light-harvesting sensitizer and rhenium bipyridine system, whereas the CO oxidation reaction was observed when a similar system was coupled with n-type NiO.<sup>70</sup> There are two main groups of metal oxides with a closed-shell electronic configuration that have been at the centre of interest for a  $CO_2$ PR system. The first group includes octahedrally coordinated  $d^0$  transition metal ions ( $Ti^{4+}$ ,  $Zr^{4+}$ ,  $Nb^{5+}$ ,  $Ta^{5+}$ ,  $V^{5+}$ , and  $W^{6+}$ ). Apart from  $TiO_2$ , which is the most prominent member of this group, other binary oxides (e.g.,  $ZrO_2$ ,  $Nb_2O_5$ , and  $Ta_2O_5$ ) have been used in  $CO_2$ PR. A number of more complex oxides referred to as titanates, niobates, tantalates, *etc.*<sup>71,72</sup> are often found in a perovskite composite,  $AMO_3$  (A = electropositive cation and M = transition metal; e.g.,  $SrTiO_3$  and  $NaNbO_3$ ), or in perovskite-related structures. Since



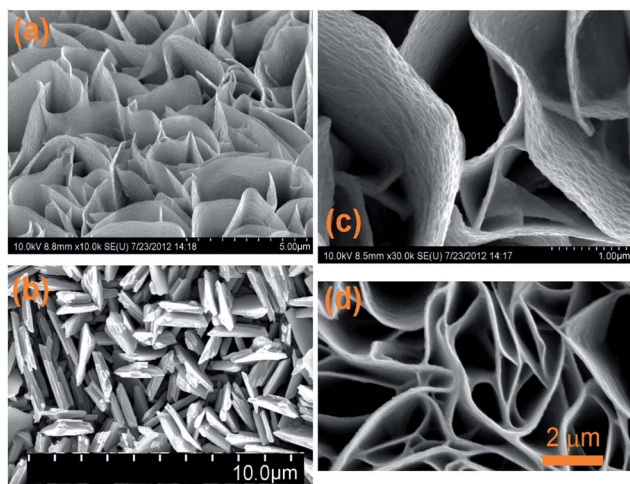


Fig. 4 SEM images of the copper sulphide nanowalls oriented vertically to the copper foil anodized at 1.5 V at room temperature (a and c), and 3.0 V (b) and 1.5 V (d) at 5 °C. Reproduced from ref. 60 with permission.

a recent published review has covered the use of perovskite oxide nanomaterials for CO<sub>2</sub> photoreduction,<sup>73</sup> this area will not be further discussed here. The second group includes main group metal oxides in a d<sup>10</sup> configuration with a general formula of M<sub>y</sub>O<sub>z</sub> or A<sub>x</sub>M<sub>y</sub>O, where M represents Ga, Ge, In, Sn, or Sb. Many of these photocatalytically active binary and ternary oxides initially found application in photocatalytic water splitting, but they have very recently started to be utilised for CO<sub>2</sub>PR.<sup>73</sup>

**Zinc-based materials.** ZnO has been widely used in the photodegradation of organic dyes and chemicals due to its direct and wide bandgap (3.37 eV).<sup>103</sup> Additionally, the bandgap and photocatalytic mechanism of ZnO are similar to those of TiO<sub>2</sub>, and thus, ZnO was also used for CO<sub>2</sub>PR. To compare the CO<sub>2</sub> photoreduction efficiency of ZnO with that of other commonly used wide bandgap semiconductors, Yahaya *et al.* employed commercially available TiO<sub>2</sub>, ZnO and NiO as photocatalysts for CO<sub>2</sub> photoreduction under 355 nm UV laser irradiation.<sup>74</sup> Among the samples, ZnO and NiO produced high yields of methanol (325 and 388 μmol g<sup>-1</sup> h<sup>-1</sup> over 1.5 h, respectively, Table 1, entries 7 and 8); whereas TiO<sub>2</sub> had the lowest production yield. In order to enhance the light absorption of commercial ZnO in the UV-vis region, ZnO was calcined to 350 °C and the ZnO obtained was immobilised onto a stainless-steel mesh to reduce the agglomeration of the photocatalyst.<sup>104</sup> The maximum conversion of CO<sub>2</sub> achieved was 11.9% (*i.e.*, percentage of CH<sub>4</sub> produced from CO<sub>2</sub> in the presence of a CH<sub>4</sub> reductant). A study revealed that the microstructure of ZnO played a more vital role than doping of ZnO with nitrogen (N-ZnO), even though the latter showed enhanced light absorption from 400 to 650 nm.<sup>75</sup> A fluffy mesoporous structured ZnO with a surface area of 29.7 m<sup>2</sup> g<sup>-1</sup> exhibited enhanced CO production (0.73 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup>, Table 1, entry 9) under UV-vis light irradiation when compared to the N-doped ZnO (1.2 m<sup>2</sup> g<sup>-1</sup>, 0.04 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup>, Table 1, entry 10). A separate publication reported that ZnO plates with

porous rectangular and assembled hexagonal morphologies (15.5 m<sup>2</sup> g<sup>-1</sup>, which was about 2.6 times that of the commercial ZnO, Fig. 5) could be obtained by using NH<sub>4</sub>Zn<sub>3</sub>(OH)<sub>6</sub>NO<sub>3</sub>.<sup>76</sup> The synthesized ZnO plates exhibited a much higher production of CO and CH<sub>4</sub> (76.35 and 20.52 ppm h<sup>-1</sup>, respectively) than the pristine hexagonal ZnO plates (44.68 and 1.57 ppm h<sup>-1</sup> of CO and CH<sub>4</sub>, respectively).

Doping has been widely used to extend the light absorption of wide bandgap semiconductors to a longer wavelength region by introducing intra-band states above the valence band. However, this approach tends to increase the recombination rate and decrease the charge mobility of the semiconductor, as discussed in Section 1. To avoid these drawbacks, the introduction of foreign cations into the binary semiconductor was considered instead of doping. For example, the ternary Zn<sub>2</sub>GeO<sub>4</sub> semiconductor was used for CO<sub>2</sub>PR under UV-vis irradiation. By fabricating ultralong and ultrathin single crystal Zn<sub>2</sub>GeO<sub>4</sub> nanoribbons, the photocatalytic reduction rate of CO<sub>2</sub> into CH<sub>4</sub> was greatly enhanced to 25.0 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup> when compared to that of the bulk Zn<sub>2</sub>GeO<sub>4</sub> (trace amounts, Table 1, entry 12).<sup>77</sup> The enhanced photocatalytic efficiency was attributed to the superb crystal quality and higher surface area (28.3 m<sup>2</sup> g<sup>-1</sup>) when compared to the bulk Zn<sub>2</sub>GeO<sub>4</sub> (0.75 m<sup>2</sup> g<sup>-1</sup>), resulting in enhanced separation of photogenerated electron-hole pairs and charge mobility. In the following year, the same group proposed the synthesis of the single crystal Zn<sub>2</sub>GeO<sub>4</sub> at 40 °C to optimise the surface area.<sup>78</sup> As a result, the surface area of the synthesized Zn<sub>2</sub>GeO<sub>4</sub> nanorods was 33.1 m<sup>2</sup> g<sup>-1</sup> which yielded 179 and 35 ppm g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup> of CO and CH<sub>4</sub>, respectively. Further increasing the temperature to 100 °C, however, decreased the surface area to 14.8 m<sup>2</sup> g<sup>-1</sup>, yielding only 3.2 and 0.4 ppm h<sup>-1</sup> of CO and CH<sub>4</sub>, respectively. By reducing the concentration of the Ge-precursor and the solvothermal time employed in the first study in 2010 (refer to ref. 77) by half, a sheaf-like superstructured Zn<sub>2</sub>GeO<sub>4</sub> was obtained and reported by the same group in 2012 (Fig. 6).<sup>79</sup> Although the CO<sub>2</sub>PR of the superstructured Zn<sub>2</sub>GeO<sub>4</sub> was not reported in this study, the optimised RuO<sub>2</sub> and Pt co-loaded Zn<sub>1.7</sub>GeN<sub>1.8</sub>O nano-sheaves after nitridation (32.33 m<sup>2</sup> g<sup>-1</sup>) could produce CH<sub>4</sub> with an apparent quantum yield of 0.024% at 450 nm (Table 1 entry 14).

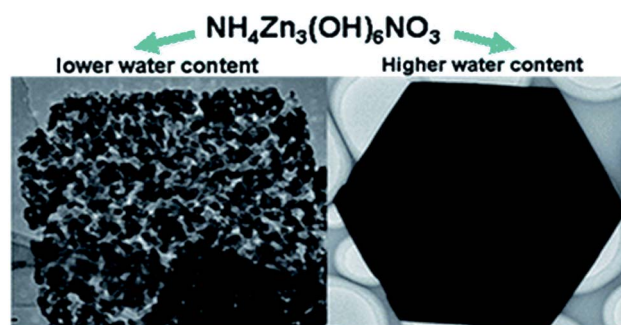


Fig. 5 ZnO synthesized from NH<sub>4</sub>Zn<sub>3</sub>(OH)<sub>6</sub>NO<sub>3</sub> with <3 mL (left) and >3 mL (right) of water. Reproduced from ref. 76 with permission.



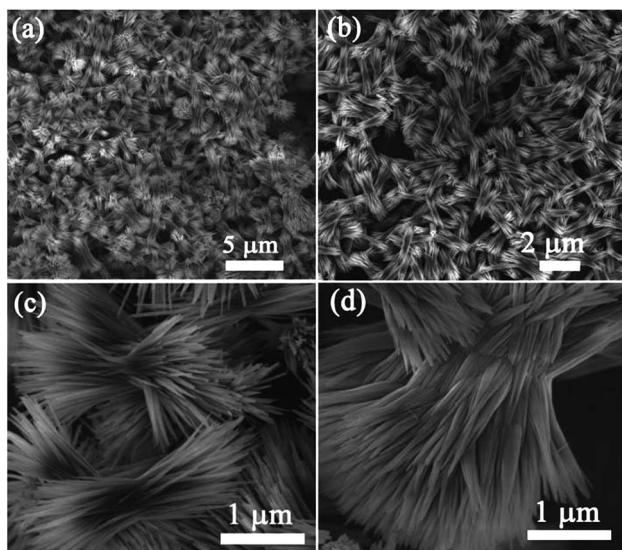


Fig. 6 SEM images of a sheaf-like  $\text{Zn}_2\text{GeO}_4$  superstructure at different magnifications. Reproduced from ref. 79 with permission.

Other nanostructured ternary Zn-based oxides were also proposed by the same group more recently, including  $\text{ZnGa}_2\text{O}$  nanosheet-scaffolded microspheres<sup>80</sup> and hexagonal nanoplate-textured  $\text{Zn}_2\text{SnO}_4$  with micro-octahedron architecture for  $\text{CO}_2$ PR application.<sup>81</sup> The unique architecture of the synthesized  $\text{ZnGa}_2\text{O}$  and  $\text{Zn}_2\text{SnO}_4$  significantly enhanced the separation of photogenerated electron-hole pairs, increased the surface area and extended light absorption. Hence, the methane yield obtained from the  $\text{CO}_2$ PR was greatly improved from trace amounts to 69 and 47 ppm  $\text{g}_{\text{catalyst}}^{-1} \text{h}^{-1}$  for the nanostructured  $\text{ZnGa}_2\text{O}$  and  $\text{Zn}_2\text{SnO}_4$ , respectively (Table 1, entries 15 and 16).

To promote co-adsorption of  $\text{CO}_2$  and  $\text{H}_2\text{O}$ , Guo *et al.* fabricated  $\text{ZnFe}_2\text{O}_4$  spinels doped with Ce.<sup>82</sup> By using *in situ* FTIR, it was found that the  $\text{CO}_2$  amount adsorbed on the surface of Ce-doped  $\text{ZnFe}_2\text{O}_4$  was much higher than that on pristine  $\text{ZnFe}_2\text{O}_4$ . This phenomenon was attributed to the increase of basicity due to the presence of alkaline  $\text{CeO}_2$  and electron density on the surface of the Ce-doped  $\text{ZnFe}_2\text{O}_4$ , thereby increasing the number of adsorption bonds between the  $\text{CO}_2$  molecules and the surface of the photocatalyst, and activating the O=C bond (Fig. 7). The formation of active  $\text{b-CO}_3^{2-}$  and  $\text{b-HCO}_3^-$  species, which could be readily translated to highly valuable products in the  $\text{CO}_2$  photoreduction, was detected. Recently, Xiao *et al.* discovered that ultrafine  $\text{ZnFe}_2\text{O}_4$  nanoparticles with a high specific surface area ( $112.9 \text{ m}^2 \text{ g}^{-1}$ ) could promote the selectivity of the photoproduction of  $\text{CH}_3\text{CHO}$  over  $\text{CH}_3\text{CH}_2\text{OH}$ , and they produced 57.8 and  $13.7 \mu\text{mol g}^{-1} \text{h}^{-1}$ , respectively, under visible light irradiation ( $>400 \text{ nm}$ ).<sup>105</sup>

**Tungsten-based materials.** Among the first group of metal oxides,  $\text{WO}_3$  has the smallest bandgap energy of 2.7 eV and as the edge of its conduction band is located at 0 V vs. NHE at pH 7, it cannot reduce  $\text{CO}_2$ .<sup>4,106</sup> However, Xie *et al.* found that when the architecture of the  $\text{WO}_3$  changed from quasi-cubic with an equal percentage of  $\{002\}$ ,  $\{200\}$  and  $\{020\}$  facets to rectangular sheet-like with dominant  $\{002\}$  facets, it could induce  $\text{CO}_2$ PR in

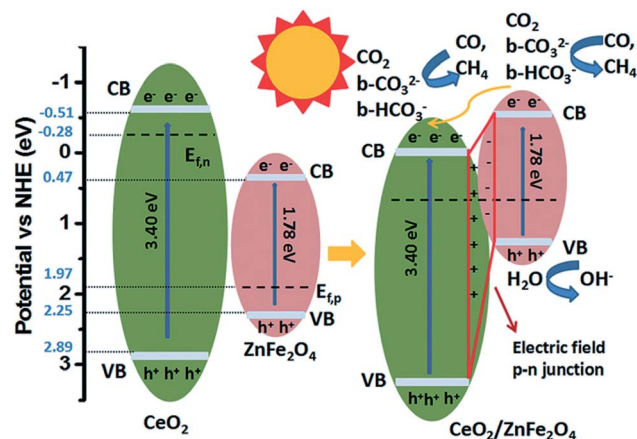


Fig. 7 Schematic of the mechanism of  $\text{CO}_2$  photoreduction with  $\text{H}_2\text{O}$  vapour over  $\text{CeO}_2/\text{ZnFe}_2\text{O}_4$ . Reproduced from ref. 82 with permission.

the presence of  $\text{H}_2\text{O}$ .<sup>83</sup> The change of the predominantly exposed facets had a significant effect on the electronic configuration of  $\text{WO}_3$ , in which the rectangular sheet-like  $\text{WO}_3$  possessed a slightly larger bandgap (2.79 eV) and its conduction band was increased by 0.3 eV. As a result, the conduction band was positioned slightly above the  $\text{CH}_4/\text{CO}_2$  potential ( $-0.24 \text{ V}$ ), inducing the methane formation from  $\text{CO}_2$  at a rate of  $\sim 0.34 \mu\text{mol g}_{\text{catalyst}}^{-1} \text{h}^{-1}$ . Chen *et al.* found that the conduction band of  $\text{WO}_3$  was shifted to a more negative position ( $-0.42 \text{ V}$ , bandgap energy: 2.79 eV) with a stronger reducing driving force, when the ultrathin ( $\sim 4\text{--}5 \text{ nm}$ ) single crystal  $\text{WO}_3$  was synthesized using a solid-liquid phase arc discharge route in an aqueous solution.<sup>84</sup> The yield obtained from the  $\text{CO}_2$ PR was  $\sim 1.1 \mu\text{mol g}_{\text{catalyst}}^{-1} \text{h}^{-1}$  under visible light irradiation ( $\lambda > 420 \text{ nm}$ ), whereas commercial  $\text{WO}_3$  powder produced only trace amounts of methane under the same conditions. A high aspect ratio of ultrathin  $\text{W}_{18}\text{O}_{49}$  exhibited extended optical properties in the visible and near infrared regions due to the presence of a large amount of oxygen vacancies (Fig. 8).<sup>85</sup> The synthesized

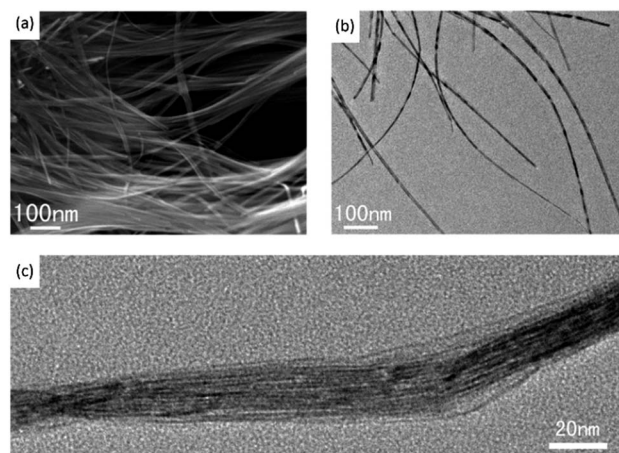


Fig. 8 SEM (a), TEM (b) and high resolution TEM (c) images of  $\text{W}_{18}\text{O}_{49}$ . Reproduced from ref. 85 with permission.



ultrathin  $W_{18}O_{49}$  exhibited the photoreduction of  $CO_2$  to  $CH_4$  at  $70^\circ C$  under visible light irradiation without a co-catalyst (Fig. 8, Table 1 entry 20). The study observed that the selectivity towards  $CH_4$  over other hydrocarbons (*e.g.*, ethanol and acetone) was as high as 95%.

The introduction of foreign elements into tungsten oxide, which generated ternary  $Bi_2WO_6$ , was reported.<sup>86</sup> The  $Bi_2WO_6$  with predominant  $\{001\}$  facets was proposed to be the most energetically favoured reactive surface for  $CO_2$  dissociation, resulting in  $1.1 \mu mol g_{catalyst}^{-1} h^{-1}$  of methane under visible light irradiation ( $\lambda > 420 nm$ ), whereas the bulk  $Bi_2WO_6$  prepared through a solid state reaction produced only trace amounts of methane. Cheng *et al.* also proposed that the microstructure of  $Bi_2WO_6$  could enhance  $CO_2$  adsorption.<sup>87</sup> A template-free anion exchange strategy was used to synthesize hollow microspheres of  $Bi_2WO_6$  (Fig. 9a and b). The synthesized  $Bi_2WO_6$  exhibited higher  $CO_2$  adsorption capacity when compared to  $BiVO_4$  and  $BiPO_4$  nanoparticles without hollow structures (Fig. 9c and d, respectively), leading to high photo-conversion of  $CO_2$  into methanol.

**Niobate-based materials.** Niobates with a perovskite structure have gained some attention because they share many characteristics (*i.e.*, non-toxicity, stability, and indirect wide bandgap) with titanates. Moreover, the conduction band of niobates is slightly more reductive than that of titanates, suggesting that niobates could be a more suitable material for  $CO_2$ PR. A study had shown that the microstructure of  $NaNbO_3$  played an important role in its photocatalytic activity.  $NaNbO_3$  nanowires ( $653 ppm h^{-1} g^{-1}$ , Fig. 10a) with a smaller bandgap (3.2 eV) and larger surface area ( $12.0 m^2 g^{-1}$ ) exhibited much higher methane formation from  $CO_2$  when compared to the  $NaNbO_3$  bulk (3.2 eV,  $1.4 m^2 g^{-1}$ ,  $22 ppm h^{-1} g^{-1}$ , Fig. 10b).<sup>88</sup> The enhanced photocatalytic activity was proposed to be due to the high crystallinity, high aspect ratio and anisotropic effect of the synthesized nanowires. Li *et al.* also demonstrated that the size and microstructure of photocatalysts play an important role in their photocatalytic activity.<sup>89</sup>  $KNb_3O_8$  and  $HNb_3O_8$

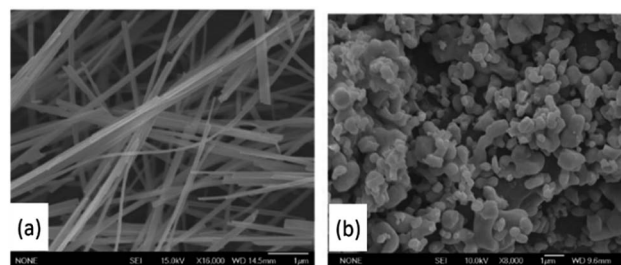


Fig. 10  $NaNbO_3$  nanowires (a) and bulk particles (b). Reproduced from ref. 88 with permission.

nanobelt samples with a higher surface area, which were  $28.8$  and  $39.4 m^2 g^{-1}$ , respectively, exhibited a  $\geq 10$  times higher photoproduction rate of methane from  $CO_2$  than the irregularly shaped  $KNb_3O_8$  and  $HNb_3O_8$  samples, which were  $2.7$  and  $6.5 m^2 g^{-1}$ , respectively (Table 1 entry 24 and 25). A similar observation was reported by Xie *et al.*, in which the nanoplates of  $SrNb_2O_6$  with an increased surface area revealed improved chemisorption of  $CO_2$  and the separation of photogenerated electron-hole pairs.<sup>107</sup> As a result, more products, such as  $CO$  and  $CH_4$ , were obtained from the  $CO_2$ PR compared to the  $SrNb_2O_6$  nanorods and nanoparticles with a lower surface area. A more recent study revealed that the nanorod-structured  $SrNb_2O_6$  ( $1.78 m^2 g^{-1}$ ,  $51.2 \mu mol g_{catalyst}^{-1} h^{-1}$ ) exhibited a higher photoreduction rate and selectivity towards  $CO$  evolution over  $H_2$  ( $>95\%$ ) than the  $SrNb_2O_7$  nanoflakes ( $3.85 m^2 g^{-1}$ ,  $6 \mu mol g_{catalyst}^{-1} h^{-1}$ ,  $\sim 39\%$ ) and  $SrNb_2O_6$  nanoparticles even though the latter possessed a higher surface area.<sup>90</sup> This phenomenon was attributed to the separation of the reduction and oxidation sites on the nanorods that decreased the recombination of photogenerated electron-hole pairs.

**Tantalum-based materials.** Tantalum-based semiconductors have been widely used as a photocatalyst for water splitting. Having higher potentials than  $TiO_2$  and above the reduction potential of  $CO_2/CH_3OH$ ,  $H_2CO_3/CH_3OH$ ,  $HCO_3^-/CH_3OH$  and  $CO_3^{2-}/CH_3OH$ ,  $Ta_2O_5$  can be employed for the  $CO_2$ PR.<sup>92</sup> However, the large bandgap energy of  $Ta_2O_5$  ( $\sim 3.9 eV$ ) has restricted its light absorption in the visible region.<sup>108</sup> Hence, Sato *et al.* used  $N-Ta_2O_5$  to couple a series of ruthenium bipyridine catalysts for the photocatalytic  $CO_2$  reduction to formic acid under visible light irradiation ( $405 nm$ ).<sup>109</sup> The production rate was found to be  $\sim 70 \mu mol g_{catalyst}^{-1} h^{-1}$  in an acetonitrile/triethanolamine mixture. To enhance the electron transportation and suppress the electron-hole recombination,  $Ta_2O_5$  was immobilised on reduced graphene and  $NiO_x$  was used as the co-catalyst (Fig. 11).<sup>91</sup> The highest photoproduction rate of methanol from 3%  $NiO_x$ - $Ta_2O_5$ -1% reduced graphene sample was  $\sim 20.83$  and  $197.92 \mu mol g_{catalyst}^{-1} h^{-1}$  without and with the presence of  $NaHCO_3$  solution, respectively.

Much effort has focused on  $InTaO_4$  as the ternary tantalum-based semiconducting photocatalyst for  $CO_2$ PR. For instance, Pan and Chen demonstrated that  $InTaO_4$  could reduce  $CO_2$  to methanol ( $\sim 1.1 \mu mol g_{catalyst}^{-1} h^{-1}$ ) in  $0.2 M KHCO_3$  aqueous solution under visible light irradiation.<sup>110</sup> The methanol production rate could marginally increase to  $1.2 \mu mol g_{catalyst}^{-1}$

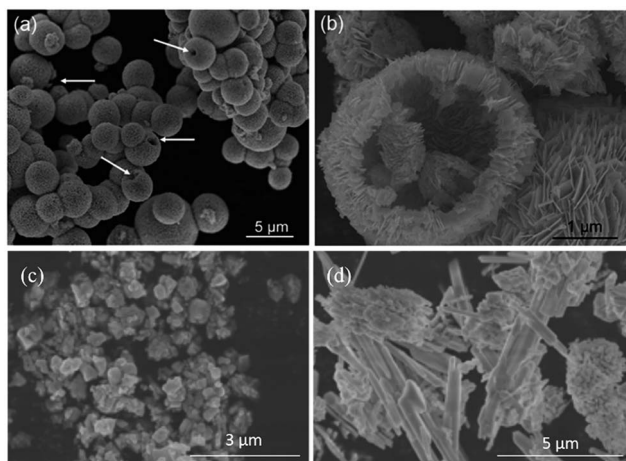


Fig. 9 SEM images of  $Bi_2WO_6$  hollow microspheres (a and b),  $BiVO_4$  (c) and  $BiPO_4$  (d) reproduced from ref. 87 with permission.



$\text{h}^{-1}$  after coupling 1 wt% NiO as the co-catalyst and  $1.4 \mu\text{mol g}_{\text{catalyst}}^{-1} \text{h}^{-1}$  after the application of reduction–oxidation pre-treatment. Tuning the size and crystallinity of  $\text{InTaO}_4$  nanoparticles resulted in the bandgap energy range from 2.6 to 3.0 eV and could also enhance the production of methanol from  $\text{CO}_2$ .<sup>111</sup> The highest production rate was about  $2.7 \mu\text{mol g}_{\text{catalyst}}^{-1} \text{h}^{-1}$  when 1.0 wt% NiO was added as the co-catalyst. The methanol generation from  $\text{InTaO}_4$  was further enhanced by introducing core–shell Ni/NiO nanoparticles on nitrogen doped  $\text{InTaO}_4$ , leading to  $160 \mu\text{mol g}_{\text{catalyst}}^{-1} \text{h}^{-1}$  under the irradiation of light with wavelengths ranging from 390 to 770 nm.<sup>92</sup>  $\text{KTaO}_3$  was also used to reduce  $\text{CO}_2$  to CO under visible light irradiation.<sup>112</sup> Three samples were synthesized with different bandgaps ranging from 3.5 to 3.7 eV and yielded the highest amount of CO at  $\sim 62 \text{ ppm g}_{\text{catalyst}}^{-1} \text{h}^{-1}$ .

Recently,  $\text{LaTa}_7\text{O}_{19}$  and  $\text{CaTa}_4\text{O}_{11}$  (bandgap energies of 4.1 and 4.5 eV, respectively) were shown to be active for  $\text{CO}_2\text{PR}$ . CO was produced after loading with 1 wt% Ag co-catalyst due to the preferable conduction band positions ( $50$  and  $70 \mu\text{mol g}_{\text{catalyst}}^{-1} \text{h}^{-1}$ , respectively, in the presence of 0.1 M  $\text{NaHCO}_3$  under the irradiation of a 400 W high-pressure mercury lamp).<sup>93</sup>  $\text{NaTaO}_3$  doped with different elements, such as Mg, Ca, Sr, Ba and La, has been proposed as a highly active photocatalyst for  $\text{CO}_2\text{PR}$  using water as the electron donor in the presence of a Ag co-catalyst under UV irradiation.<sup>94</sup> Among the samples, Ba-doped  $\text{NaTaO}_3$  loaded with 1.0 wt% Ag co-catalyst using the liquid-phase reduction method exhibited the highest CO production and selectivity from  $\text{CO}_2$  ( $\sim 50 \mu\text{mol g}_{\text{catalyst}}^{-1} \text{h}^{-1}$  and 56%, respectively).

Quaternary tantalates have been developed recently and revealed to be active for  $\text{CO}_2$  photoreduction in the presence of water.<sup>113</sup>  $\text{K}_2\text{RETa}_5\text{O}_{15}$  (RE = rare-earth element, namely La, Ce, Pr, Nd, Y, and Sm) was fabricated using the flux method with KCl, which favoured the rod-like morphology, followed by calcination treatment at  $1150 \text{ }^\circ\text{C}$  for 6 h.<sup>95</sup> Among the quaternary tantalates,  $\text{K}_2\text{CeTa}_5\text{O}_{15}$  possessed the smallest bandgap energy (2.42 eV,  $0.7 \mu\text{mol g}_{\text{catalyst}}^{-1} \text{h}^{-1}$ ), but  $\text{K}_2\text{YT}_5\text{O}_{15}$  photo-produced the highest amount of CO (3.86 eV,  $91.9 \mu\text{mol g}_{\text{catalyst}}^{-1} \text{h}^{-1}$ ). The addition of Y was shown to be beneficial for capturing  $\text{CO}_2$  and subsequently for photoreduction. Meanwhile, the presence of K in the tantalates played an important role in determining the growth orientation of the rod-like structure, thereby affecting the activity in  $\text{CO}_2$  photoreduction.

**Miscellaneous.**  $\text{CeO}_2$  is a basic metal oxide that can transform inert linear  $\text{CO}_2$  to  $\text{b-HCO}_3^-$  and  $\text{b-CO}_3^{2-}$  to reduce the reductive potential of  $\text{CO}_2$ .<sup>114</sup> Hence, it has recently attracted a lot of attention. However, it suffers from rapid recombination of photogenerated electron–hole pairs and possesses a large bandgap (3.0–3.4 eV), which restricts the light absorption in the UV range.<sup>115</sup> To improve the performance of  $\text{CeO}_2$  for the photocatalytic reduction of  $\text{CO}_2$  under visible irradiation, Xiong *et al.* proposed the coupling of  $\text{Ag}/\text{Ag}_3\text{PO}_4$  with  $\text{CeO}_2$  to construct heterojunctions for enhancing the separation of photogenerated electron–hole pairs and improve light absorption because  $\text{Ag}_3\text{PO}_4$  has a narrow bandgap of 2.42 eV.<sup>116</sup> The highest  $\text{CH}_3\text{OH}$  and  $\text{C}_2\text{H}_5\text{OH}$  yield obtained was 10.6 and  $7.9 \mu\text{mol g}_{\text{catalyst}}^{-1} \text{h}^{-1}$ , respectively. Zhang discovered that when Ni

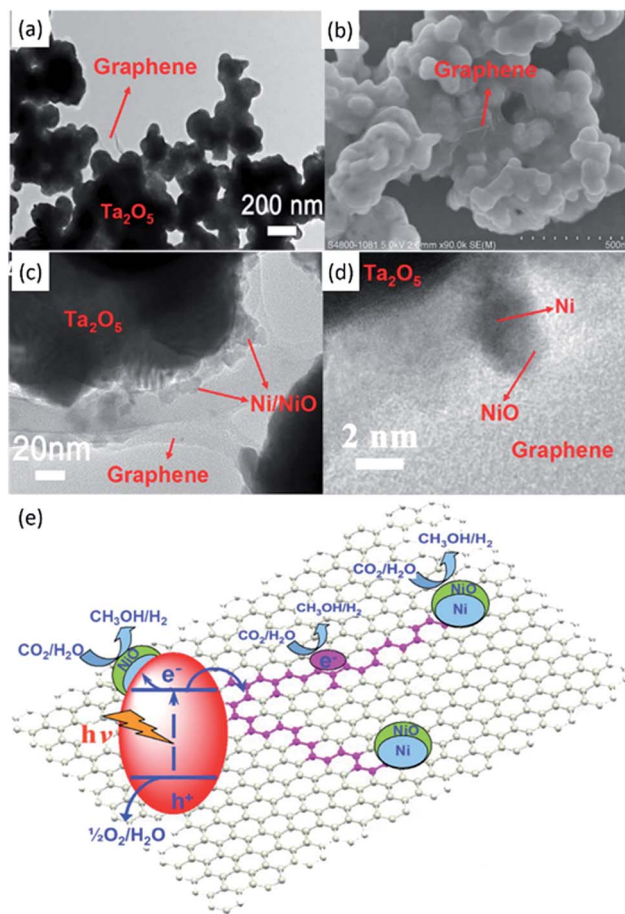


Fig. 11 TEM (a) and SEM (b) images of  $\text{Ta}_2\text{O}_5$  immobilised on reduced graphene. TEM (c) and SEM (d) images of the  $\text{NiO}_x$ - $\text{Ta}_2\text{O}_5$ -reduced graphene composite. Schematic illustration of the charge separation and transfer in the  $\text{Ta}_2\text{O}_5$ -reduced graphene system under UV-vis light (e). Reproduced from ref. 91 with permission.

was loaded on  $\text{CeO}_2$ , the nanocomposite exhibited enhanced photo(thermo)catalytic performance and inhibited carbon deposition.<sup>117</sup> Moreover, it is interesting to note that the full light spectrum response from UV to infrared of the Ni metal on  $\text{CeO}_2$  decreased the activation energy of C and CH oxidation steps, thus improving the overall photo(thermo)catalytic performance.

$\text{MgO}$  was employed to photocatalytically reduce  $\text{CO}_2$  into CO with a production rate of  $\sim 1.6 \mu\text{mol g}^{-1} \text{h}^{-1}$  over 6 h in the presence of  $\text{H}_2$  as the reductant under UV light ( $\lambda < 290 \text{ nm}$ ).<sup>118</sup> Mesoporous  $\text{Ga}_2\text{O}_3$  yielded 1.46 and  $0.21 \mu\text{mol g}^{-1} \text{h}^{-1}$  of CO and  $\text{CH}_4$ , respectively, from  $\text{CO}_2$  under visible light irradiation.<sup>119</sup> When  $\text{Ga}_2\text{O}_3$  was loaded with Ag, the photoproduction rate of CO from  $\text{CO}_2$  was  $10.5 \mu\text{mol g}^{-1} \text{h}^{-1}$  under UV light irradiation (Table 1 entry 33).<sup>96</sup> Iron oxides were proposed as a photoactive centre to induce the photocatalytic reduction of  $\text{CO}_2$ .<sup>120</sup> Using electron spin resonance spectroscopy (ESR), the photogenerated electrons from the Fe–O species were efficiently consumed by  $\text{CO}_2$  under UV irradiation.

Lamellar  $\text{BiVO}_4$  was proposed to exhibit a selective methanol production from  $\text{CO}_2$  photoreduction under visible light



irradiation.<sup>97</sup> The maximum CH<sub>3</sub>OH production rate was 5.52 μmol h<sup>-1</sup> when 0.2 g of BiVO<sub>4</sub> was suspended in 100 mL of NaOH (1.0 M) under full spectrum irradiation of a Xe lamp. The photocatalytic mechanism was proposed according to which the Bi<sup>3+</sup> sites could efficiently receive electrons from the V 3d-block bands of the BiVO<sub>4</sub> to form CO<sub>2</sub><sup>•-</sup> radical anions, leading to the formation of methoxyl radicals (·OCH<sub>3</sub>) and eventually CH<sub>3</sub>OH after hydrogen abstraction. Wang *et al.* doped the atomically thin layers of BiVO<sub>4</sub> with different percentages of Co.<sup>121</sup> The Co-doped BiVO<sub>4</sub> exhibited an efficient and stable activity for CO<sub>2</sub> photoreduction to CH<sub>4</sub>. The optimal CH<sub>4</sub> production rate was 23.8 μmol g<sup>-1</sup> h<sup>-1</sup>, which was three times higher than that of the pristine BiVO<sub>4</sub>, at 60 °C with an atmospheric CO<sub>2</sub> concentration (~400 ppm) under a UV lamp (25 W at 254 nm). The enhancement of the production rate of the Co-doped BiVO<sub>4</sub> was suggested to be due to the presence of electron enriched adsorption sites, which was contributed by the Co dopant, activating the CO<sub>2</sub> molecules for further reduction reaction.

Delafossite materials with a general stoichiometry of ABO<sub>2</sub>, in which A is a monovalent metal ion, such as Cu, Ag, and Pt, and B is a trivalent metal ion, such as Al, Ga, and Fe, as the new class of photocatalysts have also been considered for CO<sub>2</sub>PR.<sup>98</sup> CuGaO<sub>2</sub> (bandgap energy ~3.7 eV and weak absorption at 2.6 eV) and the Fe-alloyed CuGa<sub>1-x</sub>Fe<sub>x</sub>O (1.5 eV) facilitated the photo-generation of CO from CO<sub>2</sub> under the irradiation of a Xe lamp though varied amounts of Fe substituted into CuGaO<sub>2</sub> did not significantly enhance the CO<sub>2</sub> photoreduction performance (Table 1 entry 35).<sup>98</sup>

Based on the Lewis acidity of CO<sub>2</sub>, alkaline catalysts will benefit the adsorption and activation of CO<sub>2</sub>. Layered double hydroxide (LDH) materials usually possess high specific surface areas, which provide numerous active sites for the catalytic reaction. The fabricated CoAl LDH facilitated an enhanced CO<sub>2</sub> photoreduction reaction when compared to P25 due to the surface alkaline OH groups for efficient adsorption of CO<sub>2</sub> at a low concentration.<sup>99</sup> The utilisation of LDH in CO<sub>2</sub> photoreduction has been reviewed previously, and thus, interested readers may refer to the published review articles.<sup>40,122</sup>

In summary, metal oxides have shown their ability to promote photocatalytic reduction of CO<sub>2</sub>, as discussed in the previous section. However, most of these photocatalysts only work under UV irradiation due to their large bandgap energies (>3 eV). The relatively large bandgap of metal oxides originates from the valence band maximum, which is formed by O 2p orbitals and is more positive than 3 V.<sup>123</sup> Hence, if metal oxides meet the thermodynamic requirement for CO<sub>2</sub>PR and H<sub>2</sub>O photooxidation, then the bandgap of the metal oxides inevitably becomes larger than 3.0 eV, which is too wide to absorb visible light.<sup>124</sup>

### 2.3 Oxynitrides

**Tantalum-based materials.** The N 2p orbital has a higher potential energy than the O 2p orbital, which indicates that metal oxynitrides as well as metal nitrides could be employed as an efficient photocatalyst.<sup>42,125</sup> For example, in the case of

tantalum oxynitride and tantalum nitride, when nitrogen-based N 2p atomic orbitals were introduced into Ta<sub>2</sub>O<sub>5</sub>, new orbitals with a higher bound state energy are generated, resulting in a decrease of bandgap energy.<sup>42,108,126</sup> As a result, the bandgaps of TaON and Ta<sub>3</sub>N<sub>5</sub> (2.5 and 2.1 eV, respectively) are both smaller than that of Ta<sub>2</sub>O<sub>5</sub> (3.9 eV) and thus can effectively absorb visible light and drive the photocatalytic activity. Moreover, the N content plays an important role in determining the bandgap energy in oxynitrides and nitrides. Therefore, Gao *et al.* proposed a Ca-assisted urea synthesis method to controllably synthesise TaON and Ta<sub>3</sub>N<sub>5</sub> with a tailored N composition.<sup>127</sup> In addition, the initial urea : Ta molar ratio used in the proposed synthesis method was also beneficial to control the size and homogeneity of the final product. Recently, the use of the porous spherical architecture of TaON for CO<sub>2</sub> photoreduction was proposed.<sup>100</sup> The surface area of the porous spherical TaON was ~11.12 m<sup>2</sup> g<sup>-1</sup>, whereas that of commercial TaON was ~7.41 m<sup>2</sup> g<sup>-1</sup>. As a result, the CH<sub>3</sub>CHO and C<sub>2</sub>H<sub>5</sub>OH production rates from CO<sub>2</sub> using the porous TaON (0.52 and 2.03 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup>) were higher when compared to those of the commercial TaON (0.16 and 0.84 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup>) under visible light irradiation. The enhanced CO<sub>2</sub>PR in the porous TaON was attributed to the increase of surface area. In addition, the reduction of charge transfer distance and enhanced light scattering within the porous spherical structure were also suggested to play roles in enhancing the photocatalytic reduction of CO<sub>2</sub>.

The CO<sub>2</sub>PR is a multi-electron process, and a variety of products can be produced using a single semiconducting photocatalyst. The achievement of efficient and selective production of highly valuable fuels is critical for viable CO<sub>2</sub> photoreduction processes. The application of perovskite oxynitrides, such as CaTaO<sub>2</sub>N coupled with the binuclear Ru(II) complex photosensitiser and loaded with the Ag co-catalyst, revealed an enhanced selectivity for HCOOH production (>99%) from CO<sub>2</sub> under visible light irradiation due to the enhanced interfacial electron transfer.<sup>128</sup> A similar approach with the same photosensitiser and co-catalyst coupled with yttrium-tantalum oxynitride (YTON) was recently proposed by the same group.<sup>129</sup> The YTON (2.1 eV) exhibited a smaller bandgap than CaTaO<sub>2</sub>N (2.5 eV), thus extending the light absorption up to 600 nm. Moreover, the selectivity for HCOOH formation from CO<sub>2</sub> was not affected and remained as high as that in their previous study (>99%).

**Zinc-based materials.** Mesoporous ZnGeON was used as a photocatalyst for CO<sub>2</sub>PR under visible light irradiation (λ < 400 nm).<sup>130</sup> The prolonged nitridation time from 1 to 15 h decreased the Zn and O contents, in which Zn was evaporated and O was substituted by N, at 800 °C in an NH<sub>3</sub> environment. However, the crystallinity of ZnGeON was enhanced with a slight decrease in the surface area. The ZnGeO nitrided for 10 h (24.4 m<sup>2</sup> g<sup>-1</sup>) exhibited the highest CH<sub>4</sub> production rate of 2.7 ppm g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup>, which was higher than those of the ZnGeON prepared using the solid-state reaction (3.3 m<sup>2</sup> g<sup>-1</sup>, 1.1 ppm g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup>) and N-doped TiO<sub>2</sub> (2.2 ppm g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup>). As discussed in Section 2.2, zinc-based materials like ZnGa<sub>2</sub>O<sub>4</sub> possessed a bandgap energy of ~4.5 eV, in which



the valence band was mainly composed of O 2p and the conduction band was formed from the hybridization of Ga 4s and Zn 4p orbitals.<sup>80,101</sup> In order to reduce the bandgap energy, nitridation of mesoporous ZnGa<sub>2</sub>O<sub>4</sub> was proposed. The nitridation of ZnGa<sub>2</sub>O<sub>4</sub> stimulated the hybridization of Zn 3d, N 2p and O 2p and formed the valence band, whereas the conduction band was composed of the Ga 4s and 4p orbitals.<sup>101</sup> As a result, the bandgap energy was reduced to 2.5 eV, resulting from the uplifting of the maximum of the valence band and lowering of the minimum of the conduction band. To further enhance the CO<sub>2</sub>PR performance, a ZnGa<sub>2</sub>ON solid solution was modified with ZnAl<sub>2</sub>O<sub>4</sub> that acted as the CO<sub>2</sub> arrester (Fig. 12).<sup>101</sup> The increase of Zn content had also decreased the bandgap energy to 2.3 eV. Benefiting from the mesoporous structure, smaller bandgap and enhanced CO<sub>2</sub> adsorption ability, the ZnAl<sub>2</sub>O<sub>4</sub>-modified ZnGa<sub>2</sub>ON showed a methane generation rate of 9.2 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup> from CO<sub>2</sub>, which was 9 times higher than that of the pristine ZnGa<sub>2</sub>ON under visible light irradiation (λ ≥ 420 nm).

## 2.4 Nitrides

**Gallium-based materials.** Through engineering the nanostructure of the co-catalyst used, selectivity using semiconducting nitrides could be enhanced dramatically, as demonstrated by AlOtaibi *et al.*<sup>102</sup> The decoration of the non-polar GaN nanowires with the Rh core and amorphous Cr<sub>2</sub>O<sub>3</sub> shell co-catalyst significantly increased the production rate of CH<sub>4</sub> from 1.3 (bare GaN) to 3.5 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup>, but the CO production rate decreased from 1130 (bare GaN) to ~120 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup> in 24 h (Fig. 12). Due to the effective collection of photogenerated electrons by the Rh core and amorphous Cr<sub>2</sub>O<sub>3</sub> shell co-catalyst, no apparent reductive reaction (*e.g.*, photoreduction of CO<sub>2</sub> to CO) occurred on the surface of GaN without Rh coverage. As a result, the product selectivity towards CH<sub>4</sub> was enhanced in the CO<sub>2</sub>PR under UV-visible light irradiation. In addition, the decoration of the GaN nanowires with Rh/Cr<sub>2</sub>O<sub>3</sub> could suppress the back reaction that formed H<sub>2</sub>O from H<sub>2</sub> and O<sub>2</sub>, and offered adsorption sites for CO<sub>2</sub>.

In summary, both metal oxynitrides and nitrides have shown their capability to photoreduce CO<sub>2</sub> with more favourable optical properties when compared to metal oxides. Unfortunately, these groups of materials have not been extensively explored.

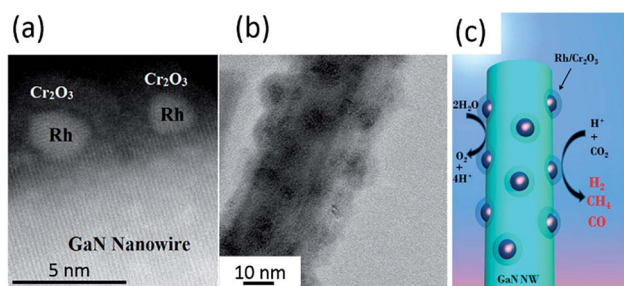


Fig. 12 (a) High-resolution TEM image and (b) TEM image of Rh/Cr<sub>2</sub>O<sub>3</sub> core/shell decorated GaN nanowires. (c) Schematic of the photoreduction processes of CO<sub>2</sub> on Rh/Cr<sub>2</sub>O<sub>3</sub>-decorated GaN nanowires. Reproduced from ref. 102 with permission.

## 3. Hole scavengers for CO<sub>2</sub> photoreduction

Various semiconducting materials have been proposed as photocatalysts for CO<sub>2</sub>PR under UV and/or visible irradiation, as discussed in Section 2 and summarised in some other references.<sup>7–10,103,131</sup> However, the quantum efficiency of the CO<sub>2</sub> photo-conversion into hydrocarbons remained low and could not rely only on the development of photocatalysts. System optimisation plays an important role in optimising the conversion rate and selectivity as well as photocatalyst stability. Therefore, the increase of CO<sub>2</sub>PR efficiency through the introduction of a hole scavenging agent has gained significant interest. In this section, the use of organic and inorganic hole scavengers is reviewed.

### 3.1 Inorganic hole scavengers

As discussed in Section 2.1, metal sulphides suffer from photocorrosion in an aqueous dispersion due to the oxidation of lattice S<sup>2-</sup> ions to elemental S and subsequently to sulphates.<sup>51</sup> Hence, the addition of reducing agents to prevent the oxidation of the lattice S<sup>2-</sup> ions by scavenging the photogenerated holes was proposed. Kanemoto *et al.* achieved a cumulative quantum yield of 72% with irradiation of UV light at 313 nm (*i.e.*, 75.1 and 1.7 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup> of HCOOH and CO, respectively) when NaH<sub>2</sub>PO<sub>2</sub> and Na<sub>2</sub>S (0.35 and 0.24 M, respectively) were added into the system that contained ZnS as the photocatalyst for CO<sub>2</sub>PR (Table 2 entry 1).<sup>132</sup>

A systematic study was recently carried out to investigate the effect of Na<sub>2</sub>S as the hole scavenger for ZnS on the CO<sub>2</sub>PR at λ = 345 nm.<sup>133</sup> The study elucidated that the photogenerated holes on the surface of ZnS were directly consumed by Na<sub>2</sub>S, whereas photogenerated electrons were pumped into the conduction band simultaneously. In addition, the behaviour of the reaction rate at different pH values resembled that of the solubility of CO<sub>2</sub>, discarding the direct participation of the HCO<sub>3</sub><sup>-</sup> and CO<sub>3</sub><sup>2-</sup> in the photoreduction process. This observation was supported by a very recent study, in which KHCO<sub>3</sub> was used as the hole scavenger in an aqueous system with ZnS.<sup>134</sup> The study demonstrated that KHCO<sub>3</sub> acted as an effective hole scavenger as well as a buffer to mitigate the pH change induced by the CO<sub>2</sub> saturation. This phenomenon, however, was not observed when only K<sub>2</sub>SO<sub>3</sub> was used as the hole scavenging agent.

The optimised solution with 0.1 g of colloidal ZnS, 0.1 M K<sub>2</sub>SO<sub>3</sub> and 0.5 M KHCO<sub>3</sub> achieved 464.2 and 81.3 μmol of HCOOH and CO, respectively, under UV-vis irradiation (Table 2 entry 2).<sup>134</sup> The selectivity towards HCOOH was reported to be 12.5%, and this could be improved to 95.0% when Cd was added to the colloidal ZnS suspension as the co-catalyst.

Inorganic salts (*e.g.*, NaOH, Na<sub>2</sub>S, *etc.*) have been reported to have a significant effect on CO<sub>2</sub>PR.<sup>97,132,134,135</sup> The addition of NaOH had been shown to increase the solubility of CO<sub>2</sub> compared to pure H<sub>2</sub>O because the OH<sup>-</sup> ions provided by NaOH in aqueous solution reacted with the dissolved CO<sub>2</sub>, and transformed into CO<sub>3</sub><sup>2-</sup> and further into HCO<sub>3</sub><sup>-</sup> in the CO<sub>2</sub> saturated system.<sup>136</sup> It was suggested that the high



Table 2 Photocatalytic CO<sub>2</sub> reduction yields obtained by various photocatalyst after adding hole scavenger(s)

| No.              | Photocatalyst   | Hole scavenger  | Function/role  | Product(s) of CO <sub>2</sub> photoreduction (μmol g <sub>catalyst</sub> <sup>-1</sup> h <sup>-1</sup> )   | Ref. |
|------------------|---|---|--|--|------|
| <b>Inorganic</b> |   |   |  |  |      |
| 1.               | ZnS quantum crystallites                                      | 0.70 M NaH <sub>2</sub> PO <sub>2</sub><br>0.48 M Na <sub>2</sub> S   | Electron donor<br>Sulphur vacancy suppressor               | HCOOH 75.10<br>CO 1.70   | 132  |
| 2.               | ZnS   | 0.1 M K <sub>2</sub> SO <sub>3</sub><br>0.1 M K <sub>2</sub> SO <sub>3</sub> <sup>+</sup><br>0.5 M KHCO <sub>3</sub>          | Hole scavenger<br>Electron donor<br>Buffer solution        | HCOOH ~250.00<br>HCOOH 580.30  | 134  |
| 3.               | Zn-doped Ga <sub>2</sub> O <sub>3</sub>                       | 0.1 M NaHCO <sub>3</sub>  | Hole scavenger   | CO 117.00  | 138  |
| 4.               | Ag-loaded SrNb <sub>2</sub> O <sub>6</sub>                    | 0.1 M NaHCO <sub>3</sub>  | CO <sub>2</sub> supply                                     | CO ~4.00   | 71   |
| 5.               | Ag-loaded Sr <sub>2</sub> Nb <sub>2</sub> O <sub>7</sub>      |   |  | CO ~38.40  |      |
| 6.               | Sr and Ag co-loaded NaTaO <sub>3</sub>                        | 0.1 M NaHCO <sub>3</sub>  | Buffer for supplying CO <sub>2</sub>                       | CO ~352.00   | 94   |
| 7.               | Ni-Al layered double hydroxides (LDHs)                        | 0.1 M NaCl  | Hole scavenger   | CO 112.80  | 139  |
| 8.               | BiVO <sub>4</sub>   | 1.0 M NaOH  | Hole scavenger   | CH <sub>3</sub> OH 5.52  | 97   |
| <b>Organic</b>   |   |   |  |  |      |
| 9.               | CdS   | Acetonitrile + dichloromethane<br><br>1.0 mol dm <sup>-3</sup> 2-propanol   | Surface modifier<br><br>Surface modifier<br>Hole scavenger | (CH <sub>3</sub> ) <sub>2</sub> CO ~0.24 μmol with 70 μM of CdS powder<br><br>(CH <sub>3</sub> ) <sub>2</sub> CO ~0.19 μmol with 70 μM of CdS powder | 140  |
| 10.              | Wurtzite-ZnS  | Isopropanol<br>Ethylene glycol  | Hole scavenger<br>Hole scavenger                           | HCOOH ~40 ppm g <sub>catalyst</sub> <sup>-1</sup> h <sup>-1</sup><br>HCOOH ~90 ppm g <sub>catalyst</sub> <sup>-1</sup> h <sup>-1</sup>               | 141  |
| 11.              | Mononuclear Ru complex/C <sub>3</sub> N <sub>4</sub>          | 4 : 1 of <i>N,N</i> -dimethylacetamide (DMA) : triethanolamine (TEOA)   | Proton quencher<br>Electron donor                          | HCOOH ~1100.00   | 142  |
| 12.              | Bi-nuclear Ru complex/Ag-loaded C <sub>3</sub> N <sub>4</sub> | 4 : 1 of <i>N,N</i> -dimethylacetamide (DMA) : triethanolamine (TEOA)   | Proton quencher<br>Electron donor                          | HCOOH ~2115.00   | 143  |
| 13.              | Bi-nuclear Ru complex/Ag-loaded C <sub>3</sub> N <sub>4</sub> | 1.0 mM ethylenediaminetetraacetic acid disodium salt dihydrate, EDTA·Na <sub>2</sub>  | Electron donor   | HCOOH ~31.67   | 144  |
| 14.              |   | 1.0 mM ethylenediaminetetraacetic acid disodium salt dihydrate (EDTA·Na <sub>2</sub> ) + 0.1 M K <sub>2</sub> CO <sub>3</sub> | Electron donor<br>Surface modifier                         | HCOOH ~83.33   |      |

concentration of HCO<sub>3</sub><sup>-</sup> present in the system could accelerate the photoreduction reaction, thereby enhancing the photoreduction performance.<sup>136,137</sup> The direct consumption of HCO<sub>3</sub><sup>2-</sup> to produce CO was observed in a CO<sub>2</sub> aqueous solution using a Zn-doped Ga<sub>2</sub>O<sub>3</sub> photocatalyst.<sup>138</sup> This observation was further investigated by Nakanishi *et al.*, in which the production of CO occurred after transforming HCO<sub>3</sub><sup>-</sup> into CO<sub>2</sub>.<sup>94</sup> Therefore, the addition of HCO<sub>3</sub><sup>-</sup> increased the CO<sub>2</sub> supply in the aqueous system, but did not enhance the numbers of reacted electrons and holes. In other words, under basic conditions, H<sub>2</sub> production could be significantly suppressed, resulting in a high selectivity of CO from CO<sub>2</sub>. Jin *et al.* concurred that the addition of NaOH promoted the formation of HCO<sub>3</sub><sup>-</sup>; however, the production of methanol was observed in the photocatalytic system with a BiVO<sub>4</sub> photocatalyst.<sup>97</sup>

The addition of NaOH, Na<sub>2</sub>CO<sub>3</sub> and NaHCO<sub>3</sub> to the CO<sub>2</sub> photoreduction aqueous solution was shown to promote the photoproduction of CO, whereas H<sub>2</sub>SO<sub>4</sub> and NaCl were found to favour water splitting, leading to the production of H<sub>2</sub>.<sup>90,94,136,137</sup> However, a recent study has proposed that the inclusion of Cl<sup>-</sup> from NaCl could scavenge the photogenerated holes for the CO<sub>2</sub> photocatalytic reduction process in a aqueous solution over Ni-

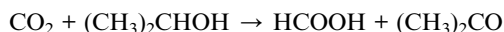
Al layered double hydroxides.<sup>139</sup> The selectivity towards CO over H<sub>2</sub> was 86% when NaCl was added into the photoreduction system (~6.95 and ~1.16 μmol g<sup>-1</sup> h<sup>-1</sup> of CO and H<sub>2</sub>, respectively), whereas in the absence of a scavenging agent, the selectivity towards CO over H<sub>2</sub> was only 54% (~3.15 and ~2.63 μmol g<sup>-1</sup> h<sup>-1</sup> of CO and H<sub>2</sub>, respectively). The authors also pointed out that the addition of NaHCO<sub>3</sub> and Na<sub>2</sub>CO<sub>3</sub> promoted the production of H<sub>2</sub> *via* the reduction of H<sup>+</sup> derived from H<sub>2</sub>O instead of the CO<sub>2</sub>PR. Neither Na<sub>2</sub>SO<sub>4</sub> nor NaNO<sub>3</sub> could positively enhance the production of CO. A similar extent of CO evolution and selectivity were also observed when other chloride salts were added, namely CsCl (~8 μmol g<sup>-1</sup> h<sup>-1</sup> of CO, 82%), MgCl<sub>2</sub> (~7 μmol g<sup>-1</sup> h<sup>-1</sup> of CO, 82%) and CaCl<sub>2</sub> (~7.5 μmol g<sup>-1</sup> h<sup>-1</sup> of CO, 82%). However, other halogenide salts (NaBr and NaI) showed a weaker photoreduction ability than NaCl.

### 3.2 Organic hole scavengers

Since the last century, ZnS had been used as photocatalyst for the CO<sub>2</sub>PR.<sup>146,147</sup> Triethylamine (TEA) showed its feasibility to be used as the hole scavenger, inhibiting the photocorrosion for sulphite photocatalysts.<sup>7</sup> In ZnS systems, 2-propanol was one of



the common hole scavengers for CO<sub>2</sub>PR. A previous report proposed that the light energy could be stored within the light-induced reaction given as



The Gibbs free energy of this reaction is +62.8 kJ mol<sup>-1</sup> at 25 °C.<sup>147</sup>

In a system of Cd-loaded ZnS, CO<sub>2</sub> photoproducted formic acid with a quantum efficiency 32.5% in the presence of 1 M 2-propanol.<sup>147</sup> Further increasing the Cd concentration resulted in the formation of CO. A study revealed that CdS was capable of photoreducing CO<sub>2</sub> to CO when *N,N*-dimethylformamide (DMF) containing 1 v/v% water was employed in the system.<sup>148</sup> A similar observation was reported, in which CO was photoproducted when CdS was dispersed in DMF under the irradiation of a 500 W mercury lamp with a 300 nm cut off filter.<sup>140</sup> When DMF was substituted with a low polarity solvent, such as CCl<sub>4</sub> and CH<sub>2</sub>Cl<sub>2</sub>, CO production was dominant, whereas when using a high polar solvent, such as H<sub>2</sub>O, formate was produced. This was because the adsorbability of the CO<sub>2</sub><sup>•-</sup>, an intermediate species after the activation of CO<sub>2</sub>, was strongly dependent on the polarity of the solvent used. For instance, low polarity molecules enabled strong adsorption of CO<sub>2</sub><sup>•-</sup> on the Cd sites of CdS through the carbon atom of CO<sub>2</sub><sup>•-</sup>, which was not highly solvated in solvents of low polarity, resulting in the formation of CO. When high polar solvents were used, CO<sub>2</sub><sup>•-</sup> was stabilised in the system and established only weak interactions with the photocatalyst. As a result, CO<sub>2</sub><sup>•-</sup> tended to react with a proton and produced formate.

A recent study suggested that the CO<sub>2</sub> photoreduction process can be greener when glycerol, which is a green solvent derived from vegetable oil, was used as the hole scavenger instead of petroleum-derived solvents.<sup>141</sup> In this study, wurtzite ZnS facilitated the photoproduction of formic acid from CO<sub>2</sub> with an apparent quantum efficiency of 3.2% and 0.9% when glycerol and 2-propanol, respectively, were employed as the hole scavenger.

Cyclohexanol was used as the hole scavenger for the CO<sub>2</sub> photoreduction under UV light irradiation.<sup>149</sup> The optimised sample exhibited the production of cyclohexyl formate and cyclohexanone (178.1 and 170.2 μmol g<sub>catalyst</sub><sup>-1</sup>, respectively) after 8 h. The authors elucidated that the production of cyclohexanone was slightly lower than that of cyclohexyl formate because some of the photogenerated holes were consumed by cyclohexanol to form cyclohexyl ether.

A recent study demonstrated that a Ru(II)-complex/C<sub>3</sub>N<sub>4</sub> nanocomposite could induce the photocatalytic CO<sub>2</sub> reduction by using a mixture of solvents (*N,N*-dimethylacetamide and DMA/TEOA).<sup>142</sup> The apparent quantum efficiency achieved was 5.7% at 400 nm (Table 2 Entry 12). In addition, the product selectivity of the Ru(II)-complex/C<sub>3</sub>N<sub>4</sub> nanocomposite could be enhanced through manipulating the solvent used (Fig. 13).<sup>145</sup>

In order to avoid using organic solvents as the medium, the mononuclear Ru(II) complex proposed in a previous study<sup>142</sup> was replaced with a binuclear Ru(II) complex coupled with Ag/C<sub>3</sub>N<sub>4</sub>

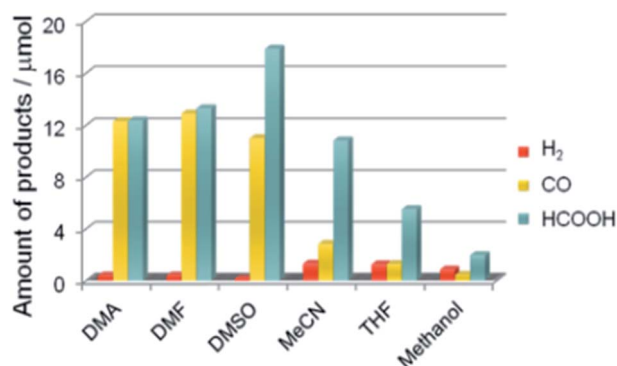


Fig. 13 Product distribution of visible light ( $\lambda > 400$  nm) CO<sub>2</sub> reduction using Ru(II) complex/C<sub>3</sub>N<sub>4</sub> as a photocatalyst and various solvents under a CO<sub>2</sub> atmosphere (DMSO: dimethyl sulfoxide, MeCN: acetonitrile, and THF: tetrahydrofuran). Reproduced from ref. 145 with permission.

and was employed as the photocatalyst (Table 2 entry 13).<sup>143</sup> Since no reduction product was obtained in pure water, a hole scavenger (ethylenediaminetetraacetic acid disodium salt dihydrate, EDTA·Na<sub>2</sub>) was added to promote the photocatalytic CO<sub>2</sub> reduction in water. The main product was HCOOH, and H<sub>2</sub> was produced as a by-product under visible light irradiation ( $\lambda > 400$  nm). Other hole scavenging agents (e.g., potassium oxalate and sodium ascorbate) were shown to be useful for the CO<sub>2</sub>PR. Among the three hole scavenging agents, sodium ascorbate exhibited the best performance with 31.7 μmol and 86% selectivity towards HCOOH. The HCOOH production could be further enhanced to 83.3 μmol with selectivity 97% when K<sub>2</sub>CO<sub>3</sub> (0.1 M) was used as an additive.<sup>144</sup> However, the production of H<sub>2</sub> was reduced by half.

The introduction of organic and inorganic hole scavenging agents has exhibited advantages to enhance the efficiency of CO<sub>2</sub>PR. The presence of hole scavenging agents in the CO<sub>2</sub>PR process is necessary if the oxidation reaction in the CO<sub>2</sub>PR cannot be inhibited by the photocatalyst. Moreover, to avoid carbon contamination and false positive errors for the photogeneration of hydrocarbons in the CO<sub>2</sub>PR process, inorganic hole scavenging agents are preferred.

## 4. Conclusions and future directions

To date, significant achievements have been made in the design and fabrication of photocatalysts and the optimisation of photocatalytic systems. CO<sub>2</sub>PR using metal sulphides, oxides, oxynitrides and nitrides accumulated so far have offered alternative photocatalytic materials other than TiO<sub>2</sub>. Material properties, including the surface area, light harvesting, and charge generation, separation and transportation, have been manipulated through the structural and morphological control during the fabrication processes, leading to enhanced CO<sub>2</sub>PR performance. Amongst the non-titania photocatalysts (metal sulphides, oxides, oxynitrides and nitrides) reviewed here, the ultrathin W<sub>18</sub>O<sub>49</sub> exhibited the highest CH<sub>4</sub> yield (2200 μmol g<sub>catalyst</sub><sup>-1</sup> h<sup>-1</sup>) from CO<sub>2</sub> under visible light irradiation. The



presence of oxygen vacancies was suggested to play an important role in the CO<sub>2</sub>PR. On the other hand, the addition of inorganic salts or organic solvents into an aqueous system has shown to effectively scavenge the photogenerated holes and/or increase CO<sub>2</sub> solubility.

Although significant studies have been carried out on CO<sub>2</sub>PR, some challenges still remain. Firstly, an in-depth understanding of the working mechanism in a CO<sub>2</sub> photoreduction process is still not well understood. Hence, a trial-and-error approach was used when fabricating photocatalysts, attempting to achieve a high CO<sub>2</sub>PR efficiency. Secondly, the insight into the CO<sub>2</sub>PR in the presence of hole scavenging agent(s) is not available. Moreover, due to this lack of knowledge, a rational design to combine state-of-art of photocatalysts with the desired hole scavenging agent(s) for carbon fuel production is difficult to achieve. Therefore, while more effort is required in material advancement, studies of the combined effect of the proposed photocatalyst with a hole scavenger should be encouraged. In addition, further investigation of CO<sub>2</sub>PR at the molecular level through *in situ* characterisation techniques should be carried out as this is key to boosting the efficiency of CO<sub>2</sub>PR.

## Conflicts of interest

There are no conflicts to declare.

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