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A comparative overview of MXenes and metal oxides as cocatalysts in clean energy production through photocatalysis

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Photocatalytic conversion of H₂O, CO₂ and N₂ represents one promising approach to harvest and store solar energy, for which efficient visible light responsive semiconductors are inevitable. Often, the presence of a small amount of an additional component called a "cocatalyst", is required to synergistically enhance the performance of the photocatalyst. Tremendous efforts were made in the past to identify inexpensive materials to be used as cocatalysts, for which metal oxides (MOs) are one of the traditional choices. Among alternative categories of materials investigated, the recently discovered MXenes display enormous potential owing to their unique 2D layered structure, tuneable composition, abundant surface functionalities and superior electronic conductivity. Specifically, MOs and MXenes encompass a variety of distinct as well as analogous characteristics that allows them to be tailored to different extents. Unfortunately, a comprehensive overview covering the synthetic, structural and photocatalytic aspects of MOs and MXenes is not available as of now. Herein, we intend to summarize the progress achieved so far in these two families of materials to be used as cocatalysts for the photoconversion of H₂O, CO₂ and N₂. Followed by a general introduction, we briefly outline the fundamental principles and the role of cocatalysts in photocatalytic reactions. A discussion regarding the use of MOs and MXenes as cocatalysts for the conversion of H₂O, CO₂ and N₂ is then provided in separate sections. Critical assessment regarding structure and morphology control, surface properties and stability concerns can not only help to recognize the challenges that limit further advancement, but can also highlight the future research directions of these materials for the effective transformation of H₂O, CO₂ and N₂.

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10th anniversary statement

J. Mater. Chem. A is dedicated to publish original contributions to research at the interface of materials science and chemistry. To celebrate the 10th anniversary of *J. Mater. Chem. A*, we devoted a special review discussing, what else than materials, new developed materials such as MXenes and old, but still unfashionable materials such as metal oxides. As you will be able to see while reading this review, the use of the two types of materials as cocatalysts leads to obtaining truly extraordinary hybrid composites with applicability in various photocatalytic reactions. Speaking about photocatalysis in general, and in particular about finding new materials with direct applications in the photocatalysis field, *J. Mater. Chem. A* was, is and it will be a landmark for both specialized and non-specialized readers passionate about novelty. Therefore, what binds us to the *J. Mater. Chem. A* is this very passion for novelty in the field of materials chemistry. Now reaching its 10th anniversary, we wish *J. Mater. Chem. A* to continue to surround us with the joy of reading outstanding papers that arouse our attention and appetite for new discoveries. Happy Anniversary, *J. Mater. Chem. A!*

1. Introduction

Being the major contributor to global warming, the emission of large quantities of CO₂ by the burning of fossil fuels demands immediate attention. The harmful effects arising from the increasing concentration of atmospheric CO₂, coupled with the

progressive depletion of the fossil fuel reserves drives the quest for alternative energy sources. Therefore, energy harvesting in a clean and sustainable manner remains the most crucial challenge the world is currently facing. Solar energy is promising in this regard and represents a readily available and exploitable substitute to fossil fuels.^{1–3} Solar energy needs to be captured and converted into useful forms, especially considering the intermittency (day/night, seasons, etc.). One such existing and established technology is electricity generation using solar cells.^{4,5} Solar electricity needs to be converted into chemical fuels, since these are the primary energy feedstock for

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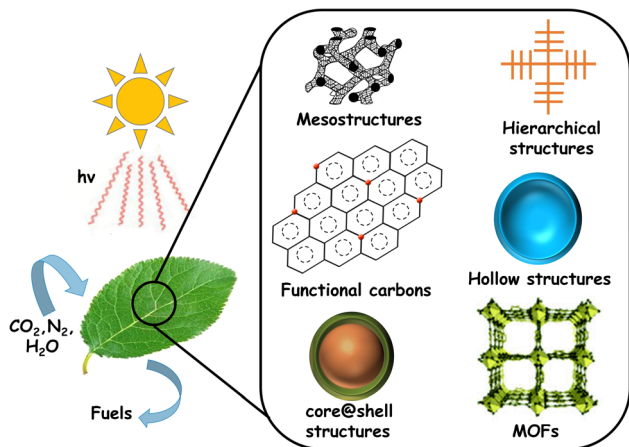


Fig. 1 Schematic representation of solar fuel production from CO_2 , H_2O and N_2 , along with the photocatalytic structural variants commonly studied.

industrial and transportation sectors. However, such transitions are difficult, demand additional costs and are time consuming. Moreover, there are technological challenges involving storage and transportation in a cost-effective and convenient manner. Therefore, direct production of fuels and chemicals using solar energy is considered a more attractive option.

Inspired by the natural process of photosynthesis, efforts were made in the recent past to create solar fuels using semiconductor photocatalysts, called ‘artificial leaves’, as shown in



Integrated in the Catalytic Material and Catalysis group, the research team is part of the National Institute of Materials Physics, Romania (NIMP) (infim.ro). Back row, left to right: Dr Ștefan Neațu (senior researcher), Dr Mahesh Nair (Postdoc), and Dr Florentina Neațu (senior researcher) and lower row, left to right: Dr Mihaela Florea (senior researcher) and Alexandra Iacoban (PhD student). The design, characterisation, and investigation of the physical properties of novel materials are the main goals of the NIMP team's research. Their primary areas of applications implies heterogeneous catalysis, energy conversion, electrochemistry, fuel cells, and materials surface science.

Fig. 1. In this case, photon energy from the sun is used to generate electron–hole pairs to drive redox reactions.^{6–10}

Most of the initial studies focused on the production of H_2 by splitting H_2O using photocatalysts.^{11,12} Apart from being an energy carrier itself, H_2 can be used to generate hydrocarbons by photoreduction of CO_2 , thereby creating a significant environmental impact by converting a greenhouse gas into a useful commodity or for N_2 photofixation toward a better solar-to-chemical energy conversion.

In short, the production and development of a technology that utilizes sunlight to efficiently convert H_2O and CO_2 into H_2 , hydrocarbons, or syngas ($\text{CO} + \text{H}_2$) can revolutionize the concept of fuel production. Here, functional materials that encompass desired photocatalytic properties such as enhanced solar light absorption, high charge separation, appropriate band gaps, *etc.*, form a major requirement.^{13–16} Compositional optimization of these materials was the earliest and most straightforward approach followed to achieve this. A variety of oxides, oxynitrides, metal organic frameworks (MOFs), carbonaceous materials, *etc.*, were studied to identify the best candidates for producing solar fuels from H_2O and CO_2 .^{17–22} Unfortunately, materials that satisfy all the practical requirements have not yet been found. On the bright side, several of these studies confirmed the dependence of photocatalytic properties on the structural features and morphology of the materials.^{13,15,17}

One proven strategy to enhance the photoactivity of a semiconductor is to integrate a small amount of a secondary component. This secondary component called a ‘cocatalyst’ facilitates the separation of charge carriers by forming an interface with the photocatalyst. Also, it can serve as the active site for the surface reaction to take place.^{23–26} Preliminary studies on the use of noble metals (*i.e.*, Pt, Pd, Ag, and Au) as cocatalysts were promising, however, with limited commercial applicability owing to their high costs.^{27–29} Numerous non-noble metal alternatives were therefore investigated as cost-effective photocatalysts. So far, transition metal-based materials including oxides, hydroxides, sulfides, *etc.*, have been widely explored.^{30–34} More recently, the introduction of 2D transition metal carbides, called MXenes, as cocatalysts was found to significantly promote photocatalytic H_2O splitting and CO_2 reduction.^{35–40}

In general, performance optimization primarily relies on structural and compositional aspects of the materials employed. Metal oxides (MOs) can accommodate a large number of cationic components in varying amounts in their structure, thereby allowing the long-range tailoring of their physicochemical properties. In conjugation, the noteworthy progress achieved so far in inorganic synthesis makes it possible to prepare these materials in a variety of nanostructures and morphologies. Similarly, MXenes, regardless of their short period of evolution, exhibited significant potential to host a number of metallic components in varying proportions in their structure.^{41–44} Several recent reports also claim the possibility of preparing morphological variants of these materials.³⁸ An additional advantage exclusive to MXenes is the



presence of inherent surface functionalities that can be useful for catalytic applications.

Several reviews exist regarding the synthesis and applications of MXenes, including photocatalysis.^{35–45} Moreover, theoretical results and insights on how MXenes and/or MOs contribute to photocatalysis as cocatalysts exist already in the literature.^{46–52} The present review summarizes the major milestones achieved regarding the use of MOs and MXenes as cocatalysts for hydrogen production through the water splitting reaction and photocatalytic CO₂ and N₂ reduction reactions. To maximize the synergistic interactions, it is crucial to maintain optimum contact between the semiconductor and cocatalyst, which depends primarily on the synthetic procedures and nanoscale structural arrangement. Therefore, we separately discuss the general advantages of nanostructures in photocatalysis, the progress so far on the development of synthetic approaches and the resultant structure evolution for MXenes and MOs. Furthermore, applications of these two categories of materials for photocatalytic H₂O, CO₂ and N₂ conversion will be summarized. Comparisons will be made highlighting the advantages and disadvantages, where appropriate. In addition, challenges and future directions for MOs and MXenes regarding the photocatalytic conversion of H₂O, CO₂ and N₂ will be provided from the perspective of the authors.

2. Fundamentals of photocatalysis and the relevance of nanostructures

In general, photocatalysis involves light absorption, charge separation (electrons and holes) and surface reactions initiated by these photogenerated charge carriers (see Fig. 2). Photon energy corresponding to the band gap of the semiconductor will be absorbed resulting in the generation of electrons (e⁻) in the conduction band and holes (h⁺) in the valence band. The reactants will be adsorbed at the active sites on the surface of the photocatalyst. The photogenerated charge carriers (e⁻ and h⁺) can get trapped, recombine, or migrate towards the surface, to drive the desired transformation of the adsorbed reactant species. The products thus formed will then be desorbed from the surface.^{53,54} During water splitting, the photogenerated holes will interact with H₂O, producing O₂ and H⁺. The H⁺ ions will consume the e⁻ residing at the conduction band forming H₂.

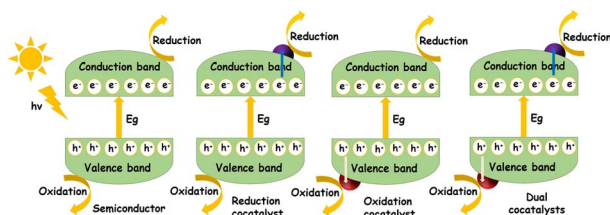
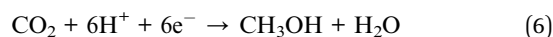
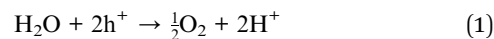


Fig. 2 Fundamental principles in photocatalysis including light absorption, charge separation, and surface reactions, along with the influence of cocatalysts.

On the other hand, photoreduction of CO₂ is more complex involving multiple electrons that can end up in a series of low molecular weight organic products. Therefore, in this later case, reaction selectivity is also a major concern.⁵⁵ The N₂ photo-fixation process involves the use of holes generated by light irradiation to oxidize the H₂O (as in the case of water splitting), while the e⁻ are used to reduce the N₂ to ammonia.⁵⁶ The photocatalytic water splitting, CO₂ reduction and N₂ photo-fixation reactions are shown in eqn (1)–(2), (3)–(7) and (8), respectively. It is to be noted here that apart from electrons and holes, other reactive intermediates are formed during photocatalysis. These reactive oxygen species are particularly useful for the photodegradation of organic pollutants and hence will not be discussed here.⁵⁷



Thermodynamically, a reduction reaction will take place if the conduction band is located at a more negative potential and an oxidation reaction requires a more positive valence band than the oxidation potential. Unfortunately, this requirement is often satisfied by materials with much wider band gaps, prohibiting their functioning in the visible region, where the major portion of solar energy is distributed. In addition, prolonged charge separation is strongly desired to minimize the electron–hole recombination that can occur during their migration towards the surface.^{53,54}

Systematic tailoring of the structure and morphology at the nanoscale can have a profound effect on the overall photocatalytic properties of materials.^{13,15,58,59} Performance enhancement in nanostructures can be attributed to an increased surface area, reduced diffusion barriers and shortened charge transport pathways. Nanomaterials in general exhibit enhanced values of specific surface area. Since only surface atoms participate in photocatalytic reactions, a high surface area implies that more atoms are available as adsorption sites for the reactants. A high surface-to-volume ratio results in an abundant and uniform distribution of accessible active sites and often promotes charge transfer across interfaces. Several studies considered nanostructuring as a way to improve the light harvesting efficiency of photocatalysts.^{13,15,60,61} A few possible effects of nanostructures on photocatalysis are schematically shown in Fig. 3.



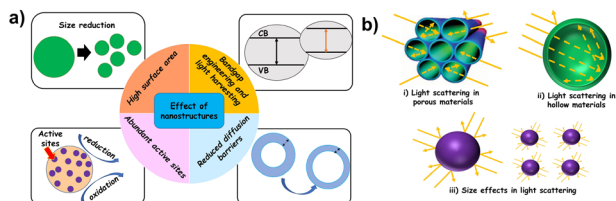


Fig. 3 (a) General influence of nanostructures in photocatalysis and (b) light scattering effects under confinement.

Nanostructures will provide an increased number of transport paths resulting in a prolonged interaction time of the photocatalyst with incoming light. This results in an enhanced surface distribution, thereby improving the light absorption effectiveness. For instance, light scattering effects observed in some nanostructures can influence the light harvesting efficiency of photocatalysts. In a conventional bulk material, surface reflection is predominant whereas in nanostructures, the incident light is scattered resulting in secondary absorptions, leading to enhanced light harvesting.^{61–63}

The effect of secondary scattering can be more pronounced where a high degree of confined spaces is available (e.g., nanoporous materials). Thin films or hollow nanostructures with thin walls could promote facile diffusion of charge carriers towards the surface adsorbed reactant species.⁶⁴ Nanoporous materials generally have particle size distribution in the micrometer regime and can prevent stability issues related to agglomeration prevalent in conventional nanosized particles. Apart from the enhancement of light absorption and transport kinetics, the quantum confinement effects inherent in nanostructures can often facilitate appropriate band gap engineering. Quantum size effects can modify the position of the valence and conduction bands in the photocatalyst. Also, confined nanoparticles can result in the generation of multiple e^-/h^+ pairs (excitons). Overall, the improved photocatalytic performance of nanostructured materials stems from a combination of their structural features, specific surface area and presence of accessible active sites.⁶⁵ In reality, photocatalytic conversion of H_2O , CO_2 and N_2 depends on the physicochemical and optical properties of the photocatalyst. Even though nanostructure optimization can improve some of these properties, it can be unfavourable to the others. Therefore, a rational approach for the design of nanostructured photocatalysts must take the disadvantages also into consideration. For instance, an increased surface energy inherent in nanomaterials can influence its thermodynamic stability. Also, high interfacial areas and short diffusion lengths in nanoscale materials can promote undesired and easy recombination of the charge carriers, influencing the reaction rates.

As mentioned above, the presence of a cocatalyst is necessary to improve the photocatalytic properties of semiconductors. In other words, the overall performance stems from the synergistic functioning of both these components in the form of a hybrid. Therefore, any discussion regarding the fundamental mechanism of photocatalysts cannot exclude the role of cocatalysts, improving a photocatalytic reaction by providing reaction sites

and by promoting charge separation. Reduction cocatalysts are used to trap electrons whereas oxidation cocatalysts are used for trapping holes. The presence of a cocatalyst should not hinder the active sites from light absorption or accessibility of the reactants and should improve the corresponding half reaction whereas dual cocatalyst-semiconductor photocatalytic systems contribute towards the advancement of the overall redox reaction.^{23–26}

3. MXenes

3.1. Short overview

The presence of a cocatalyst synergistically modifies the performance of a semiconductor photocatalyst and to maximize such interactions, it is important to maintain optimum contact. Therefore, several approaches have been developed to integrate the cocatalyst with the semiconductor. This includes the synthesis of the cocatalyst separately and then combining it with the semiconductor in a post-synthetic manner, or synthesizing the cocatalyst and the semiconductor in one pot. For instance, the synthetic procedure for MXenes is not compatible with routinely used semiconductor photocatalysts. Moreover, the thermal stability of MXenes is limited under oxidizing conditions. Therefore, post-synthetic approaches are always necessary for integrating MXenes as a cocatalyst.

3.2. Synthesis and structure evolution

Naguib *et al.*,⁶⁶ in 2011, reported a transition metal containing graphene analogue ($Ti_3C_2T_x$), called MXene, obtained by selectively extracting Al from the parent MAX phase (Ti_3AlC_2) using HF.⁶⁶ In the strategies developed so far, the synthesis of MXenes involves the selective etching of an A-layer from the MAX phase, by exploiting the differences between the metal–metal (M–A) and metal–carbon (M–X) bond strengths.^{40–42,66} The use of aqueous HF as the etchant resulted in the formation of a multilayered material with abundant –F, –O and –OH surface terminations (see Fig. 4).⁶⁷ These materials are represented by the general formula $M_{n+1}X_nT_x$, where M denotes early transition

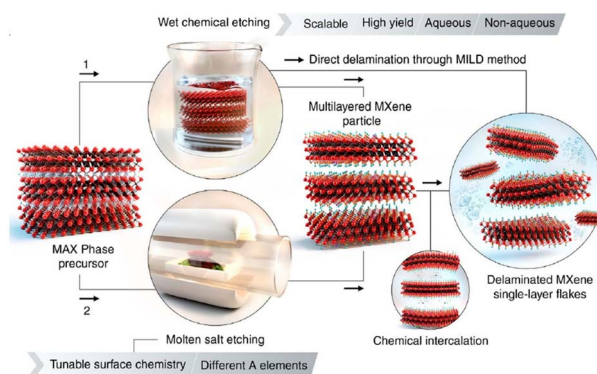


Fig. 4 Schematic representation of two etching strategies to remove the A layers from MAX phases to produce multi-layered MXenes and delaminated single layer flakes. Reproduced with permission from ref. 67, Copyright 2021 Science.



metals, X can be carbon or nitrogen, T indicates the surface functionalities such as hydroxyl groups, halides, *etc.* and n determines the number of layers, usually between 1 and 4. Recently, MXenes with $n = 5$ were discovered.⁶⁸ It is to be noted here that the etching conditions were influenced by the variations in M–Al bond strength and hence, minor disparities in etching conditions are inevitable. The presence of surface functionalities enables the post-synthetic modifications through intercalation or grafting, thereby tailoring the physicochemical properties of these materials as desired.^{69,70} With regard to the safety issues in handling HF, LiF/HCl mixture, ammonium bifluoride (NH_4HF_2), *etc.* were also found to be useful as alternatives.^{71,72} Other approaches such as electrochemical etching in HCl, hydrothermal treatment (HTT) in NaOH, iodine dissolved in anhydrous acetonitrile, molten salt such as ZnCl_2 or CuCl_2 at enhanced temperatures were also reported.^{73–77} However, in some cases, issues such as incomplete etching and the presence of impurities, could not be avoided.⁷⁴

In short, the composition of MXenes primarily depends on the parent MAX phase while the structure and surface properties are influenced by the etching protocols. The simplest version of MXenes or conventional MXenes incorporates an early transition metal attached to carbon.⁶⁶ Afterwards, more complex structural and compositional variants were discovered.^{78–83} For this, MAX phases with more than one transition metal were used as precursors. *o*-MXenes consist of multiple transition metal components with an inner layer formed by one component and the outer layer by the other. On the other hand, non-stoichiometric *i*-MXenes display an ordered array of vacancies in the metal in-plane layer.

Tremendous progress has been achieved so far in controlling the synthesis parameters to obtain 2D MXene layers. Liquid phase exfoliation of the accordion structure resulted in the expansion of interlayer space, which weakens interlayer interactions, resulting in nanosized sheets. Usually, the concept of etching was used to remove the A layer from the precursor MAX phase.

Interestingly, Ren *et al.*⁸⁵ extended this etching concept for the synthesis of porous 2D $\text{Ti}_3\text{C}_2\text{T}_x$ open structures with large surface area.⁸⁵ Delaminated colloidal solutions were partially oxidized with O_2 dissolved in the water, in the presence of Cu^{2+} as a catalyst. Selective removal of TiO_2 nanoparticles thus formed resulted in porous $\text{Ti}_3\text{C}_2\text{T}_x$ flakes. Stacking of 2D MXene sheets can limit their applicability and hence strategies are to be developed to hinder their aggregation.

Zhao *et al.*⁸⁴ proposed a hard templating method to derive hollow spheres and 3D architectures of MXene flakes and the corresponding images are provided in Fig. 5. Hollow spheres and 3D microporous films were made by using polymethyl methacrylate (PMMA) spheres as the template.⁸⁶ In a similar manner, Xiu *et al.*⁸⁷ used an ultrasound assisted aerosol spray drying method, to overcome the possible aggregation of MXene nanosheets.⁸⁷ These authors successfully assembled $\text{Ti}_3\text{C}_2\text{T}_x$ into hierarchical 3D architectures by evaporating aerosol droplets produced by ultrasonication, exhibiting geometry-based resistance to aggregation. It was also shown that by introducing polyvinyl pyrrolidone into the MXene colloidal

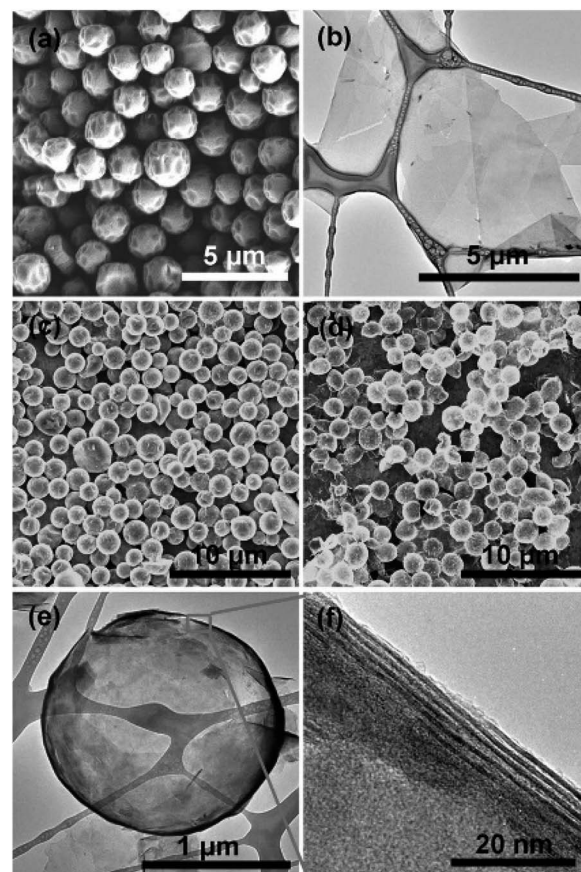


Fig. 5 (a) SEM images showing the morphology of the as-produced PMMA spheres. (b) TEM image of $\text{Ti}_3\text{C}_2\text{T}_x$ flakes. SEM image of (c) $\text{Ti}_3\text{C}_2\text{T}_x/\text{PMMA}$ hybrids and (d) hollow $\text{Ti}_3\text{C}_2\text{T}_x$ spheres after removal of PMMA, showing that the spherical shape is retained after PMMA's thermal evaporation at 450 °C. TEM images showing (e) a hollow $\text{Ti}_3\text{C}_2\text{T}_x$ sphere and (f) its wall MXene layers. Reproduced with permission from ref. 84, Copyright 2017 Wiley Publications.

solution, sphere like 3D architectures can be produced. Free standing 3D network structures were developed by assembling MXenes in the presence of templates.^{88–90} MXene quantum dots (QDs) were fabricated by dissociating their layered structure using HTT, an ultrasonic method, *etc.*^{91–94}

For photocatalytic applications, post-synthetic procedures are almost always necessary for MXenes. Simple oxidation treatment sometimes results in partial oxidation of the metallic species in MXenes, resulting in the formation of the corresponding MXene-oxide composites.^{95,96} In addition, other wet-chemical approaches including HTT, ultrasonication, *etc.* are frequently employed.^{97–99}

3.3. Photocatalytic applications for water splitting

In spite of MXenes being metallic, the surface functionalities can be effectively tailored so as to directly employ them as semiconductor photocatalysts.¹⁰⁰ Studies utilizing deformation potential theory (DPT), density functional theory (DFT) and *ab initio* molecular dynamics (AIMD) calculations clearly predicted several MXene compositions that can be used for photocatalytic



applications.^{101–104} However, experimental validation of the synthesis of such compositions and the evaluation of their photocatalytic performance are yet to be made.

3.3.1 MXenes as photocatalyst-cocatalyst precursors. Even though the direct use of MXenes as photocatalysts is not yet proven, these materials can be preferentially oxidized under controlled conditions to generate oxide particles on the surface. In this case, a cocatalyst-photocatalyst composite is found to be generated from a single MXene precursor.

For instance, by performing controlled oxidation of Nb₂CT_x using CO₂ as a mild oxidant, Su *et al.*⁹⁵ successfully synthesized Nb₂O₅/C/Nb₂C composites. The materials obtained after an oxidation time of 1 h exhibited hydrogen evolution rates 4 times higher than that of pristine Nb₂O₅, with an AQY of 0.11%. Hydrogen yield remain consistent during four consecutive cycles. The Schottky layer formed at the interface acted as an electron sink from which backward diffusion to the Nb₂O₅ photocatalyst was hindered. The presence of an amorphous carbon layer at the Nb₂O₅/C/Nb₂C interface also contributed towards the separation of photogenerated charge carriers.⁹⁵ This work confirmed that MXenes can act as precursors for photocatalysts. In another study, it was observed that carbon doped TiO₂ can be produced from Ti₃C₂T_x nanosheets under HTT.¹⁰⁵ This material exhibited a 9.7 times higher hydrogen production rate compared to commercial TiO₂ (P25) using triethanolamine (TEA) as a hole scavenger and in the absence of a cocatalyst. The authors carried out XRD analysis of the material recovered after the reaction and the pattern confirmed the structural stability of this material. This work further confirms the role of carbon in facilitating the separation of charge carriers and the use of MXene as a precursor for photocatalysts.¹⁰⁵

Li *et al.*⁹⁶ effectively controlled the surface terminations in Ti₃C₂T_x and synthesized TiO₂/Ti₃C₂T_x hybrids by performing calcination of F- terminated and OH- terminated Ti₃C₂T_x, separately.⁹⁶ OH- terminated Ti₃C₂T_x was obtained by treating Ti₃C₂T_x with NaOH. Uniformly truncated octahedral bipyramidal TiO₂ particles with exposed [001] facets were preferentially formed on the surface of F- terminated MXenes after calcination. This material exhibited two times enhanced hydrogen production compared to TiO₂/Ti₃C₂T_x produced from OH-terminated counterparts.⁹⁶

3.3.2 MXenes as cocatalysts. As of now, MXenes are being used as cocatalysts that can promote the separation of photo-generated charge carriers, thereby facilitating the photocatalytic activity of semiconductors (*e.g.*, TiO₂). Being economical and non-toxic, with a band gap ≈ 3 eV, TiO₂ is the most studied photocatalyst so far. Its band structure permits both reduction and oxidation reactions under UV light irradiation as initially shown by Fujishima and Honda.^{106,107} Table 1 summarizes several examples of MXene-based photocatalysts used in the most recent years for hydrogen production through water splitting. The synthetic approaches to combine different semiconductor photocatalysts with MXenes, and several reaction parameters are also presented. These reports demonstrated that the use of MXenes as cocatalysts could greatly improve the

photoactivity compared to the use of solely the base photocatalyst.

Wang *et al.*⁹⁷ synthesized TiO₂/Ti₃C₂T_x composites in different weight ratios (5–50% Ti₃C₂T_x) under HTT. Photocatalytic hydrogen evolution from water using methanol as a hole scavenger was monitored for these materials under visible light irradiation. Inclusion of 5 wt% Ti₃C₂T_x resulted in a 400% enhancement in hydrogen evolution compared to pure rutile TiO₂. Photocatalytic activity was further found to be improved by replacing Ti₃C₂T_x with Ti₂CT_x or Nb₂CT_x, among which 5% Nb₂CT_x exhibited the best performance among the series, with an apparent quantum yield (AQY) of 0.39%. Stability tests performed for three cycles using TiO₂/Ti₃C₂T_x exhibited consistent results.

According to the authors, the hydrothermal synthesis method allowed an intimate contact between the semiconductor and the cocatalyst surfaces resulting in a flow of photogenerated electrons. A Schottky barrier thus formed at the interface prevented the electrons from diffusing back, thereby preventing charge recombination. In addition, the highest value of the work function for Nb₂CT_x (4 eV) among those of the series reflected a larger Schottky barrier explaining its best performance among the series (see Fig. 6).⁹⁷

In a similar manner, Li *et al.*¹⁰⁸ synthesized TiO₂/Ti₃C₂T_x nanoflowers from Ti₃C₂ under HTT followed by ion exchange and calcination.¹⁰⁸ Initially, Na₂Ti₃O₇-Ti₃C₂ composites were synthesized hydrothermally, and were then treated with HCl to replace Na⁺ with H⁺ producing H₂Ti₃O₇-Ti₃C₂. These materials were then calcined at 300–500 °C to produce TiO₂/Ti₃C₂T_x. Optimum materials obtained after calcination at 500 °C displayed overall water splitting in the absence of a sacrificial agent, with an AQY of 5.86%. The better performance was attributed to the ability of 3D porous nanoflower-like morphology to provide more active sites, and its greater ability to reflect and scatter light and reduce the diffusion length of photogenerated charge carriers. The Schottky junction formed at the interface hindered charge recombination resulting in more electrons participating in H₂ evolution and more holes participating in O₂ evolution.¹⁰⁸

Su *et al.*⁹⁸ synthesized a series of Ti₃C₂T_x/TiO₂ composite photocatalysts by adding an aqueous dispersion of stacked monolayer Ti₃C₂T_x in different weight ratios to commercial TiO₂ (Degussa P25), under sonication.⁹⁸ Photocatalytic tests performed in the presence of methanol as a hole scavenger indicated a 9 times higher hydrogen evolution rate (AQY = 15.8%) for the composite with 5 wt% MXene, compared to P25. Interestingly, this composite material exhibited a 2.5 times better performance than when the multi-layer MXene counterpart was used as the cocatalyst. The significantly enhanced activity is attributed to the superior electron conductivity of monolayer Ti₃C₂T_x and enhanced charge separation at the interface. It is to be noted here that after four cycles, the hydrogen production rate decreased from 2650 $\mu\text{mol g}^{-1} \text{h}^{-1}$ to 2300 $\mu\text{mol g}^{-1} \text{h}^{-1}$. Since the structural and surface properties of the photocatalyst recovered after the reaction remained unchanged, the observed decrease in the rate was attributed to the probable detachment between TiO₂ and Ti₃C₂T_x.⁹⁸



Table 1 Compilation of synthetic and photocatalytic aspects of MXene-based materials in hydrogen production through the water splitting reaction^a

Photocatalyst/cocatalyst	Synthetic method	Hole scavenger	Light region	Optimum H ₂ production rate [$\mu\text{mol g}^{-1} \text{h}^{-1}$]	AQY [%]	Ref.
TiO ₂ /Ti ₂ CT _x	HTT (95 °C for 4 h)	Methanol (25%)	Visible	~32	0.27	97
TiO ₂ /Ti ₃ C ₂ T _x				17.8	0.15	
TiO ₂ /Nb ₂ CT _x				~46	0.39	
Nb ₂ O ₅ /C/Nb ₂ C	Oxidation under CO ₂ (850 °C for 0.5–1.5 h)	Methanol (25%)	Visible	7.81	0.11	95
TiO ₂ /Ti ₃ C ₂ nanoflowers	HTT (140 °C for 12 h), ion exchange (HCl, 24 h) and calcination in air (300–500 °C for 3 h)	Methanol (20%)	UV-visible	783.11	5.86 (350 nm)	108
TiO ₂ /Ti ₃ C ₂ T _x composites	Sonication	Methanol (25%)	UV	2650	15.8 (305 nm)	98
C-doped TiO ₂ derived from Ti ₃ C ₂ T _x	HTT (160 °C for 9 h)	TEA (10%)	Full spectrum	33.04	NP	105
Truncated octahedral bipyramidal TiO ₂ /Ti ₃ C ₂ hybrids	Calcination in air (550 °C for 4 h)	Glycerol (10%)	Full spectrum	~130	0.9 (365 nm)	96
0D/1D/2D Ag/Nb ₂ O ₅ @Nb ₂ CT _x nano hybrids	HTT (180 °C for 0.5–10 h) and <i>in situ</i> photodeposition	Methanol (7%)	Full spectrum	682	1.6 (313 nm)	109
La ₂ Ti ₂ O ₇ /Ti ₃ C ₂ composites	HTT (200 °C for 24 h)	Glycerol (7%)	UV	824	NP	110
CdS/Ti ₃ C ₂ composites	HTT (180 °C for 12 h)	Methanol (40%)	Visible	1594	NP	111
1D/2D CdS/Ti ₃ C ₂ nanocomposites	HTT (160 °C for 48 h)	Lactic acid (22%)	Visible	14 342	40.1 (420 nm)	111
1D/2D CdS/Ti ₃ C ₂ composites	HTT (160 °C for 48 h)	Lactic acid (10%)	Visible	2407	35.6 (420 nm)	112
2D/2D CdS/Ti ₃ C ₂ T _x composites	HTT (200 °C for 5 h)	TEA (20%)	Simulated solar light	63.53	2.28	113
2D/2D CdS/Ti ₃ C ₂ T _x composites	HTT (80 °C for 48 h)	Lactic acid (10%)	Visible	3226	47 (420 nm)	114
CdS/TiO ₂ -C (derived from Ti ₃ C ₂ T _x)	γ -ray irradiation	Lactic acid (20%)	Visible	1480	NP	115
CdS/Mo ₂ C	HTT 180 °C for 24 h)	Lactic acid (17%)	Visible	17 964	48.06 (420 nm)	116
2D/2D ZnIn ₂ S ₄ /2D Ti ₃ C ₂ T _x sandwich like hierarchical heterostructures	Reflux (80 °C for 2 h)	TEA (10%)	Visible	3475	11.14 (420 nm)	117
BiVO ₄ @ZnIn ₂ S ₄ /Ti ₃ C ₂ QD	Ultrasonication	None	Visible	6.16	2.9 (460 nm)	118
Ti ₃ C ₂ @TiO ₂ /ZnIn ₂ S ₄ ternary composites	HTT (180 °C for 3 h)	Na ₂ S/Na ₂ SO ₃	Simulated solar light	1185.8	NP	119
CdS-MoS ₂ -Ti ₃ C ₂ composites	HTT (160 °C for 24 h)	Na ₂ S/Na ₂ SO ₃	Visible	9679	26.7 (420 nm)	120
MoxS@TiO ₂ @Ti ₃ C ₂ composites	HTT (160 °C for 12 h)	TEA	UV-visible-near IR	10 505.8	7.535	121
CdS@Au/Ti ₃ -x C ₂ T _y ternary composites	Solvothermal treatment (180 °C for 12 h)	Lactic acid (10%)	Visible	5371	16.7 (420 nm)	122
2D/2D g-C ₃ N ₄ /Ti ₃ C ₂ nanosheets	Electrostatic self-assembly	TEA (10%)	Visible	72.3	0.81 (400 nm)	99
2D/0D g-C ₃ N ₄ /Ti ₃ C ₂ composites	Freeze drying	TEA (15%)	Simulated solar light	5111.8	3.654	123
p-g-C ₃ N ₄ /Ti ₃ C ₂ T _x hollow spheres	Electrostatic self-assembly with a sacrificial template	TEA (10%)	Visible	982.8	NP	124
2D/3D g-C ₃ N ₄ /Ti ₃ C ₂ hybrid	Calcination under N ₂ (550 °C for 2 h)	TEA (10%)	Visible	116.2	NP	125
0D black phosphorous/2D Ti ₃ C ₂ /2D g-C ₃ N ₄	Grinding and sonication	TEA (10%)	Visible	18 420	17.6 (420 nm)	126
ChI/Ti ₃ C ₂ T _x	Wet chemical	Ascorbic acid	Visible	1010	NP	127
ChI-1@ChI-2/0Ti ₃ C ₂ T _x	Wet chemical	Ascorbic acid	Visible	143	NP	128

^a ~ Data taken from a graphical representation of the corresponding reference; NP: value not provided; TEA = triethanolamine.

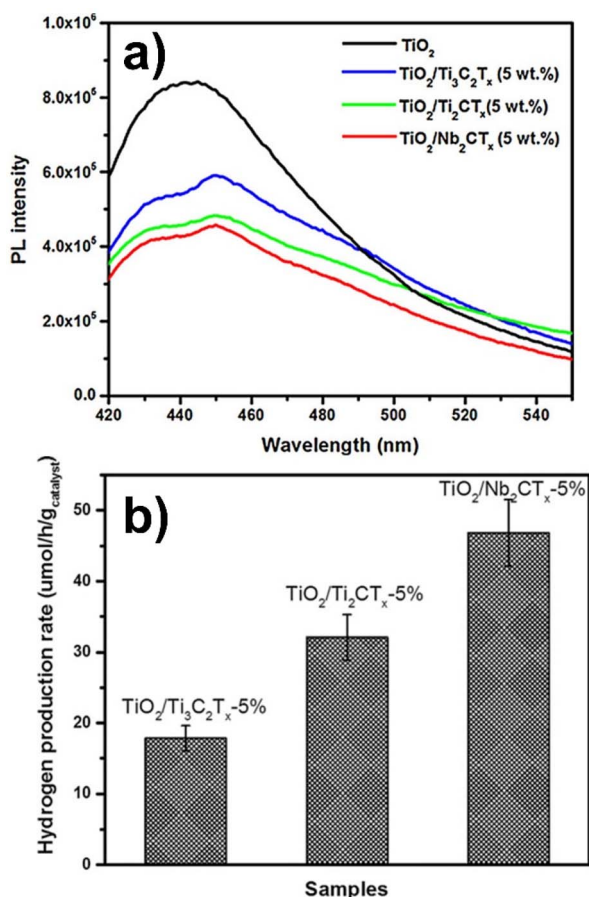


Fig. 6 (a) Photoluminescence spectra of pure TiO₂ along with those of the materials with different 5 wt% metal carbide cocatalysts. The excitation wavelength was 400 nm. (b) Photocatalytic hydrogen production rates obtained for different metal carbide cocatalysts. Reproduced with permission from ref. 97, Copyright 2016 European Chemical Societies Publishing.

In an attempt to study the charge carrier dynamics, Debow *et al.*¹²⁹ observed that band bending resulted in the formation of a Schottky barrier at the contact point between TiO₂ and Ti₃C₂T_x. The hot electrons generated from MXenes are rapidly (180 fs) transferred into the conduction band of TiO₂, indicating strong electronic coupling.¹²⁹

Peng *et al.*¹⁰⁹ fabricated 1D Nb₂O₅ nanorod arrays with a [001] orientation on 2D Nb₂CT_x under HTT followed by the photo-deposition of Ag nanoparticles. While the low work function of -OH terminated Nb₂CT_x promoted the trapping of photo-generated holes from Nb₂O₅ nanorods, Ag nanoparticles acted both as an electron reservoir and active sites for hydrogen evolution, resulting in an effective spatial separation of charge carriers. Under solar light irradiation, these nanohybrids produced 824.2 μmol g⁻¹ h⁻¹ using glycerol as the sacrificial agent.¹⁰⁹ Studies regarding stability and reusability were also carried out by these authors. It was found that hydrogen production remained unchanged after three sequential cycles. Moreover, the morphology, phase composition and surface

functional groups of Ag/Nb₂O₅@Nb₂CT_x recovered after the reaction, was not altered significantly.

In another interesting study, Wang *et al.*¹¹⁰ hydrothermally synthesized La₂Ti₂O₇/Ti₃C₂T_x composites. The introduction of 20 wt% Ti₃C₂T_x resulted in a 16 times higher hydrogen production rate than that of pristine La₂Ti₂O₇, in the presence of methanol. Solid-state NMR and Raman spectroscopy clearly indicated the formation of graphene quantum dots in Ti₃C₂T_x. In contrast to the vast majority of reports on MXene-based photocatalysts, these authors proposed that graphene quantum dots produced *in situ* during the HF etching process, rather than Ti₃C₂T_x itself, played the role of a co-catalyst during the photocatalytic reaction, by suppressing the charge recombination.¹¹⁰ However, in our understanding, no further reports have appeared in this regard.

Metal sulfides are another category of semiconductors widely investigated for photocatalytic applications. In comparison to MOs, these materials comprise a narrow band gap, enabling a visible light response, and hence, several metal sulfides were also coupled with MXenes in photocatalytic water splitting.

Ran *et al.*¹¹¹ integrated Ti₃C₂T_x nanoparticles with a series of metal sulfides in different proportions. Combination with CdS resulted in the best hydrogen yields with an apparent quantum efficiency of 40.1%. Similarly, 386% and 217% enhancement of the photocatalytic performance was respectively observed for Zn_{0.8}Cd_{0.2}S and ZnS by incorporating 1% of Ti₃C₂T_x. The authors attributed this to a favourable Fermi level position and superior electronic conductivity of MXenes. Significant variations in the performance of CdS/Ti₃C₂T_x were not observed during seven successive cycling tests. A comparison between the fresh and used materials also indicated consistency in the crystalline phase, morphology and size.¹¹¹

Hydrothermally synthesized 1D CdS nanorod/2D Ti₃C₂T_x nanosheet heterojunctions displayed 7 times increased hydrogen production compared to pristine CdS nanorods with an AQY of 35.6%, under visible light irradiation in the presence of lactic acid. In this study also, no obvious decrease in hydrogen production was observed for five consecutive cycles. The enhanced performance was attributed to the accelerated charge separation facilitated by the Schottky heterojunction formed, as well as to the synergistic interactions between the components.¹¹²

Another study highlighting the effectiveness of combining CdS with Ti₃C₂T_x under HTT, reported hydrogen evolution rates higher than that of pure CdS and CdS@Pt, with an AQY of 2.28%. The authors also noted 87.61% retention of performance after three cycles. Apart from a slightly weakened intensity of the XRD peaks, no significant changes were observed for the material recovered after the reaction.¹¹³ CdS/Ti₃C₂T_x composites forming 2D/2D Schottky heterojunctions exhibited a 5-fold enhancement of photocatalytic performance of pure CdS, with an AQY of 47%. Stable performance was observed for five consecutive cycles. In this case also, the superior performance was attributed to an intimate interface promoting rapid charge transfer and hindering their recombination, which was confirmed by DFT calculations.¹¹⁴



In another interesting study, $\text{Ti}_3\text{C}_2\text{T}_x/\text{CdS}$ nanocomposites prepared under γ -ray irradiation, upon NaOH treatment resulted in the formation of $\text{TiO}_2\text{-C}/\text{CdS}$ nanocomposites. This material exhibited hydrogen evolution rates 6.4 and 7.0 times greater than that of $\text{Ti}_3\text{C}_2\text{T}_x/\text{CdS}$ and CdS , respectively. In this case, oxygen vacancies in amorphous TiO_2 also acted as electron trapping sites to facilitate the transfer of photoexcited electrons. No significant loss in photocatalytic performance was observed for five cycles.¹¹⁵

Jin *et al.*¹¹⁶ synthesized hybrid structures of 2D Mo_2C sheets covered with CdS nanocrystals. This hybrid photocatalyst with 2.5 wt% of Mo_2C exhibited a 11% higher hydrogen production rate than noble-metal catalysts with an apparent quantum yield of 48.06%.¹¹⁶

Zuo *et al.*¹¹⁷ used $\text{Ti}_3\text{C}_2\text{T}_x$ nanosheets to support ultrathin ZnIn_2S_4 nanosheets, forming sandwich-like hierarchical heterostructures. The resultant materials exhibited improved specific surface area, pore diameter, and hydrophilicity.

A 6.6 times improvement in photocatalytic hydrogen evolution was observed for pure ZnIn_2S_4 with an AQY of 11.14%, which was attributed to facile charge transfer and inhibition of their recombination (see Fig. 7). The values of hydrogen evolved remained nearly constant after six cycles.¹¹⁷

A hierarchical core@shell structure consisting of BiVO_4 @- ZnIn_2S_4 over which Ti_3C_2 quantum dots were uniformly distributed, was obtained by following a multi-step hydrothermal approach. This material exhibited overall water splitting under visible light irradiation. Stable photocatalytic performance was observed during five cycles. The addition of Ti_3C_2 significantly improved the light absorption efficiency and range, extending towards the near infrared region. The close contact between BiVO_4 and ZnIn_2S_4 resulted in effective charge separation. MXenes captured electrons from the conduction band of ZnIn_2S_4 and thereby suppressed charge recombination. The inherent confinement effects of Ti_3C_2 in the form of quantum dots is also found to contribute.¹¹⁸

Similarly, Huang *et al.*¹¹⁹ hydrothermally generated TiO_2 nanoparticles on the surface of Ti_3C_2 nanosheets, which were

then subjected to an additional HTT to obtain hierarchical Ti_3C_2 @ TiO_2 / ZnIn_2S_4 structures. Under simulated solar light irradiation, hydrogen generation rates 9.1 times higher than that of Ti_3C_2 @ TiO_2 and 4.6 times higher than that of pure ZnIn_2S_4 were obtained. This was attributed to the superior light harvesting, availability of sufficient active sites, intimate interfacial contact and efficient charge separation resulting from the formed heterojunction. A minor decrease in H_2 amounts was observed by the authors after four cycles and was attributed to the mass loss that occurred during photocatalyst separation and recovery.¹¹⁹ The $\text{CdS-MoS}_2\text{-Ti}_3\text{C}_2\text{T}_x$ composite exhibited a 251.3% enhancement of the hydrogen generation rate of CdS-MoS_2 , under visible light irradiation, with an AQY of 26.7%. Here, the authors identified three mechanisms for the migration of photogenerated electrons from the conduction band of CdS directly to MoS_2 or $\text{Ti}_3\text{C}_2\text{T}_x$ and to MoS_2 through $\text{Ti}_3\text{C}_2\text{T}_x$. The adsorbed H^+ will get converted into hydrogen either on the surface of $\text{Ti}_3\text{C}_2\text{T}_x$ or MoS_2 edges, implying the significance of the synergy between the components. In this study, the authors noted that adding MXenes above 2% can induce agglomeration and can act as recombination sites for the charge carriers.¹²⁰ Hydrothermally enabled *in situ* growth of TiO_2 on Ti_3C_2 nanosheets followed by the distribution of MoS_2 on the surface resulted in the formation of $\text{Mo}_x\text{S}@/\text{TiO}_2@/\text{Ti}_3\text{C}_2$ with molybdenum vacancies and double cocatalysts. This unique composite displayed a 193 times higher hydrogen production rate compared to pure TiO_2 . The authors observed no significant decrease in hydrogen evolution rates for 3 cycles. Molybdenum vacancies acted as active sites and suppressed charge carrier recombination, whereas MXenes enhanced the electron conductivity.¹²¹

Li *et al.*¹²² synthesized a ternary $\text{CdS}@/\text{Au}/\text{Ti}_{3-x}\text{C}_2\text{T}_y$ composite by a solvothermal method. In particular, core-shell structured $\text{CdS}@/\text{Au}$ nanojunctions were supported on MXenes. The composite with 1 wt% $\text{Ti}_{3-x}\text{C}_2\text{T}_y$ and 0.1 wt% Au resulted in 26.6 times enhancement of hydrogen production compared to pristine CdS . The dual Schottky barriers formed at the interface of CdS/Au and $\text{CdS}/\text{Ti}_{3-x}\text{C}_2\text{T}_y$ facilitated photogenerated electron migration from CdS while limiting back transfer, as evident from XPS analysis, Kelvin probe measurements and DFT calculations. Nanoparticles on the surface of $\text{Ti}_{3-x}\text{C}_2\text{T}_y$ nanosheets exhibited signs of aggregation after 4 h which became more pronounced after 29 h, resulting in the decline of photoactivity.¹²²

Polymeric compounds are also investigated for photocatalytic reactions among which graphitic carbon nitrides ($\text{g-C}_3\text{N}_4$) are the most promising. A proper band edge position and narrow bandgap (≈ 2.9 eV) permits this material to function under visible light. Su *et al.*⁹⁹ adapted an electrostatic self-assembly approach to synthesize 2D/2D $\text{Ti}_3\text{C}_2/\text{g-C}_3\text{N}_4$ composites by introducing varying amounts of monolayer Ti_3C_2 . Incorporating 3 wt% Ti_3C_2 resulted in a 10-fold increase in photocatalytic hydrogen evolution performance of pure $\text{g-C}_3\text{N}_4$, under visible light irradiation in the presence of TEA as a hole scavenger. The hydrogen evolution rates were consistent for 3 consecutive cycles of photocatalytic reaction. Moreover, no changes were observed in the composition and structure after

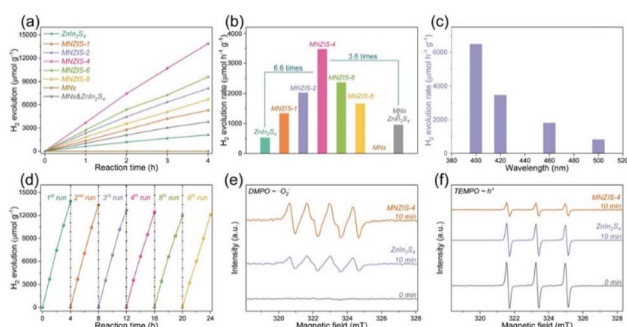


Fig. 7 (a) Time–yield plots and (b) comparison of photocatalytic H_2 evolution for ZnIn_2S_4 , $\text{Ti}_3\text{C}_2\text{T}_x$ - ZnIn_2S_4 (MNZIS-X), MNs, and a physical mixture of MNs & ZnIn_2S_4 , (c) H_2 evolution rates for MNZIS-4 with different long-pass cut-off filters, (d) cyclic experiments of H_2 evolution using MNZIS-4, and EPR spectra of (e) $\text{DMPO} \cdot \text{O}_2^-$ and (f) $\text{TEMPO} \cdot^+$ for MNZIS-4 and ZnIn_2S_4 . Reproduced with permission from ref. 117, Copyright 2020 Wiley Publications.



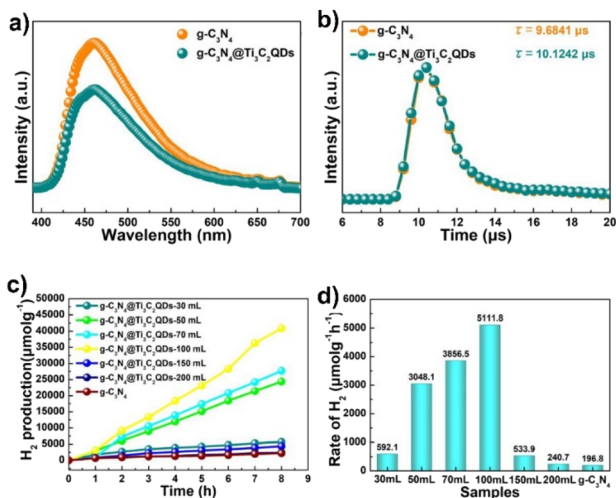


Fig. 8 (a) Steady photoluminescence (PL) and (b) time-resolved fluorescence decay spectra of $g\text{-C}_3\text{N}_4$ and $g\text{-C}_3\text{N}_4@Ti_3C_2$ QDs-100 mL composites. $\lambda_{ex} = 325$ nm. (c) Photocatalytic H_2 evolution and (d) rate of $g\text{-C}_3\text{N}_4$ NSs, Pt/ $g\text{-C}_3\text{N}_4$, Ti_3C_2 MXene sheet/ $g\text{-C}_3\text{N}_4$ and $g\text{-C}_3\text{N}_4@Ti_3C_2$ QDs. Reproduced with permission from ref. 123, Copyright 2019 The Royal Society of Chemistry.

the reaction. A slight decrease in light absorption was attributed to the probable separation of Ti_3C_2 flakes from the $g\text{-C}_3\text{N}_4$ nanosheets. The Schottky junction formed at the interface with the maximum contact surface greatly enhanced electron transport and inhibited their recombination with holes.⁹⁹

Introduction of 5.5 wt% of Ti_3C_2 QDs in $g\text{-C}_3\text{N}_4$ by the freeze-drying method resulted in 26 times higher than that of pure $g\text{-C}_3\text{N}_4$, in the presence of TEA (see Fig. 8). Interestingly, these materials also outperformed Pt/ $g\text{-C}_3\text{N}_4$. The authors attributed the better performance of these materials to the close contact between Ti_3C_2 and $g\text{-C}_3\text{N}_4$, large specific surface area and small particle size of Ti_3C_2 quantum dots. The authors also examined the recyclability and stability of these materials. No significant decrease in the amount of H_2 produced was observed during three consecutive eight hour cycles.¹²³

Kang *et al.*¹²⁴ synthesized $g\text{-C}_3\text{N}_4/Ti_3C_2T_x$ hollow spheres, by electrostatic layer-by-layer assembly, in the presence of amino-functionalized polystyrene beads as sacrificial templates. Protonation of $g\text{-C}_3\text{N}_4$ sheets was carried out using HCl. The resultant hybrid materials exhibited a 3.5 times better hydrogen production rate compared to pure protonated $g\text{-C}_3\text{N}_4$, and a 1.22-fold higher hydrogen production rate than its non-spherical counterpart. The rate of hydrogen production remained approximately the same during three repeated cycles. The 3D hollow structure enabled enhanced light absorption, while the 2D heterostructure between the components shortens the electron migration distance and high surface area providing sufficient active sites.¹²⁴

2D/3D $g\text{-C}_3\text{N}_4/Ti_3C_2$ hybrids were fabricated by calcination of melamine together with pre-formed MXenes. This unique material displayed a 6 times higher hydrogen production rate compared to pristine $g\text{-C}_3\text{N}_4$ under visible light irradiation, which was attributed to the Schottky junction formed,

preventing charge recombination, together with high electron conductivity of Ti_3C_2 and an intimate interface.¹²⁵ The performance remained stable for five cycles. Also, the crystal structure of fresh and recycled samples remained the same, indicating better stability.

Introduction of Ti_3C_2 into a heterojunction formed between black phosphorus quantum dots and $g\text{-C}_3\text{N}_4$ resulted in hydrogen evolution rates 47.2 and 19.4 times, respectively, higher than that of bulk $g\text{-C}_3\text{N}_4$ and ultrathin $g\text{-C}_3\text{N}_4$. Stable photocatalytic hydrogen evolution was observed for six consecutive cycles. With the apparent quantum yield reaching 17.6%, the composite photocatalysts with an optimum mass ratio exhibited enhanced light absorption and strong interfacial contact. Here, Ti_3C_2 acted as a bridge to accelerate charge transfer between black phosphorus and $g\text{-C}_3\text{N}_4$.¹²⁶

Li *et al.*¹²⁷ fabricated organic-inorganic composites by depositing supramolecular aggregates of a chlorophyll derivative (zinc methyl 3-devinyl-3-hydroxymethyl-pyropheophorbide a (Chl)) on the surface of $Ti_3C_2T_x$. In this case, Chl served as an organic semiconductor and MXene, a co-catalyst. For 2 wt% Chl, $52 \pm 5 \mu\text{mol h}^{-1} \text{g}^{-1}$ hydrogen evolution was observed under visible light irradiation in the presence of ascorbic acid. Efficient light harvesting followed by exciton transfer in Chl aggregates and the resultant charge separation at the interface were responsible for the observed performance.¹²⁷ In a similar study, Li *et al.*¹²⁸ grafted chlorophyll derivatives, Chl-1 and Chl-2, on the surface of 2D $Ti_3C_2T_x$ forming a Chl-1@Chl-2@ $Ti_3C_2T_x$ composite. Under visible light irradiation, the hydrogen evolution rate was found to be $143 \mu\text{mol h}^{-1} \text{g}^{-1}$ which was substantially higher than the values obtained individually for Chl-1@ $Ti_3C_2T_x$ or Chl-2@ $Ti_3C_2T_x$.¹²⁸

3.4. Photocatalytic applications for CO₂ reduction

Photocatalytic conversion of CO_2 into consumable chemicals and fuels is an ideal approach to harvest solar energy and at the same time counter the challenges related to increasing CO_2 emissions. A summary of frequently used reactants, catalysts and products is schematically shown in Fig. 9. Apart from the undesired charge carrier recombination generally encountered in photocatalysis, poor surface adsorption and activation of CO_2 molecules must also be addressed in this case. Even though addition of cocatalysts is found to be effective, noble metal free compositions are still scarce in this regard. In comparison to water splitting, fewer studies have been carried out so far using $Ti_3C_2T_x$ for the photocatalytic conversion of CO_2 . It needs to be noted here that $Ti_3C_2T_x$ (or MXenes in general) itself is a source of carbon, and therefore, additional care should be taken while interpreting the results.

Table 2 summarizes several examples of MXene-based photocatalysts used in the most recent years for CO_2 reduction. The synthetic approaches to combine different semiconductor photocatalysts with MXenes and several reaction parameters are also presented. These reports demonstrated that the use of MXenes as a cocatalyst greatly improves the photocatalytic activity compared to the use of solely the semiconductor photocatalyst. As shown in Table 2, $Ti_3C_2T_x$, as a co-catalyst,



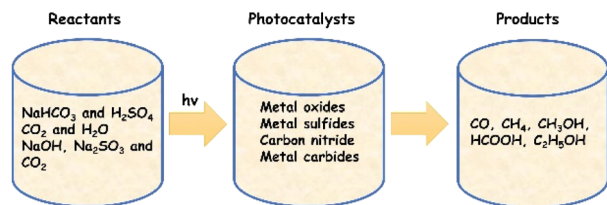


Fig. 9 Schematic representation of photocatalytic CO₂ conversion highlighting frequently used reactants, photocatalysts and possible products that can be formed.

promotes the separation of charge carriers, leading to the formation of mainly C1 products (*i.e.*, CO, formaldehyde, formic acid, methanol and CH₄). The absence of C2 product formation (*e.g.*, ethanol, ethylene or oxalic acid) indicates that Ti₃C₂T_x cannot change the energy barrier of the photocatalyst.

3.4.1 MXenes as a photocatalyst-cocatalyst precursor. Low *et al.*¹³⁰ studied the effect of calcination temperature on the stability of Ti₃C₂T_x and observed an *in situ* growth of surface TiO₂ nanoparticles. Photocatalytic CO₂ reduction was carried out over these materials under light irradiation from a Xe arc lamp. NaHCO₃ and HCl were used as the reactants to *in situ* generate CO₂ and H₂O vapours. Formation of CH₄ as a major product with small amounts of methanol and ethanol was observed for the material obtained after calcination at 550 °C. The CH₄ production rate was 3.7 times higher than that of commercial TiO₂ (P25). The labelling of the products confirmed that the hydrocarbon products originated from CO₂. Moreover, the rate of CH₄ production remained stable for five consecutive cycles. The *in situ* growth of TiO₂ resulted in an intimate contact with Ti₃C₂, facilitating the transfer and accumulation of photogenerated electrons for multi-electron CO₂ conversion to take place. In addition, increased specific surface area and enhanced light absorption properties of the composite also contributed towards performance enhancement. The authors also proposed that an excess of Ti₃C₂T_x will compete with the light absorption properties of TiO₂, thereby decreasing the efficiency.¹³⁰

In the same direction, Song *et al.*¹³² synthesized 3D hierarchical Ti₃C₂T_x and performed calcination in air to generate TiO₂ on the surface of nanosheets resulting in the formation of a heterostructure with bimodal macro-mesoporosity. These materials were used for photocatalytic conversion of gas phase CO₂ produced from the reaction between NaHCO₃ and H₂SO₄ under simulated solar light. In the absence of CO₂, product formation was not observed. Also in this case, the composites obtained after calcination at 500 °C, exhibited a 2-fold increase in the CH₄ evolution rate compared to commercial TiO₂. In this case, the 3D hierarchical structure promoted enhanced light absorption and efficient electron-hole separation whereas strong adsorption of CO₂ on Ti₃C₂ ensured the availability of electrons for the reaction.¹³²

3.4.2 MXenes as cocatalysts. Surface alkalization was found to have a profound effect on electronic conductivity of MXenes. A simple KOH treatment can effectively replace the surface -F groups in MXenes with hydroxyl groups. For the photoreduction of gas phase CO₂ carried out with H₂O as the

hydrogen source, the presence of alkalized Ti₃C₂T_x as a cocatalyst improved the CH₄ production rate of the commercial TiO₂ photocatalyst by 277 times, as shown in Fig. 10. However, on addition of above 5 wt% Ti₃C₂, P25 was shielded from light irradiation, decreasing the activity. In this study, no carbonaceous products were determined during the control experiments performed over pure Ti₃C₂T_x or by replacing CO₂ with Ar. CO₂ molecules are acidic in nature and the presence of basic OH surface functionalities on Ti₃C₂ acted as active sites for adsorption. The electron rich environment created in Ti₃C₂ after capturing the conduction band electrons from TiO₂, readily improves the reduction of the adsorbed CO₂ molecules, thus explaining the enhanced activity.¹³¹ Similar to the studies carried out by Ye *et al.*,¹³¹ surface-alkalized Ti₃C₂ was integrated with ZnO by Li *et al.* following an electrostatic self-assembly strategy. Here, the presence of 7.5 wt% of Ti₃C₂ in the composite resulted in a 35 times higher CH₄ production rate and 7 times higher CO production rate, compared to those of pure ZnO. Here, both CH₄ and CO evolution remained constant for three cycles.¹³⁸

The presence of Ru along with TiO₂ on the surface of Ti₃CN MXenes produced a mixture of CO, CH₄ and other hydrocarbons from CO₂ using water as the hydrogen source. With 0.5 wt% Ru, CO and CH₄ production rates increased by 20.5 and 9.3 times, respectively, compared to that of P25. Isotopic labelling studies confirmed CO₂ as the origin of the reaction products. Gas production rates exhibited a 13.4% decrease during the first three cycles. This was attributed to the accumulation of intermediate products on the active sites, hindering further reaction. To confirm this, the photocatalyst was recovered and washed with deionized water. This resulted in a performance similar to that observed during the first cycle.¹³³

Wang *et al.*¹³⁴ prepared a core@shell structured *meso*-TiO₂@ZnIn₂S₄ and different amounts of Ti₃C₂ were deposited on the ZnIn₂S₄ shell. This ternary photocatalyst, under UV-visible light irradiation, resulted in CO and CH₄ evolution with slightly better CH₄ selectivity. The better performance in comparison with that of the individual components was attributed to accelerated electron transfer to Ti₃C₂ as well as the enhanced CO₂ adsorption capacity of the 3D mesostructure.¹³⁴

Zeng *et al.*¹³⁵ developed a hierarchical heterostructure by decorating Cu₂O nanowires with Ti₃C₂ quantum dots (QDs) on a Cu mesh, adapting an electrostatic self-assembly strategy. Photocatalytic transformation of a CO₂ saturated H₂O medium was carried out under solar light irradiation, and selective formation of CH₃OH was observed. Product yields were found to be 8.25 times higher compared to that of Cu₂O nanowires on a Cu mesh. Interestingly, the presence of MXenes as QDs resulted in a 2.15 times higher methanol yield compared to MXene sheets with the same composition. 89% retention of methanol yield was observed after six cycles of photocatalytic reaction. The authors reported a multi-fold influence of Ti₃C₂T_x on the performance of Cu₂O NWs by enhancing charge transport, carrier density, and light adsorption, as well as by decreasing the band bending edge and limiting charge carrier recombination.¹³⁵ Taking into account the good synergy with Cu₂O, a series of Cu₂O/Ti₃C₂T_x heterostructures with varying



Table 2 Compilation of synthetic and photocatalytic aspects of MXene-based materials in CO₂ conversion^a

Photocatalyst composite	Synthetic method	Reactants	Light region	Products [$\mu\text{mol g}^{-1} \text{h}^{-1}$]	AQY [%]	Ref.
TiO ₂ /Ti ₃ C ₂	Calcination in air (350–650 °C)	NaHCO ₃ and HCl	Simulated solar light	CH ₄ (0.22) CH ₃ OH (NP)	NP	130
Alkalinized TiO ₂ /Ti ₃ C ₂	KOH treatment and physical mixing	CO ₂ and H ₂ O	Simulated solar light	C ₂ H ₅ OH (NP) CH ₄ (16.61)	1.61 0.32	131
3D hierarchical TiO ₂ /Ti ₃ C ₂ T _x heterojunction	Calcination in air (300–700 °C for 1 h)	NaHCO ₃ and H ₂ SO ₄	Simulated solar light	CO (11.74) CH ₄ (4.41)	NP	132
Ru–Ti ₃ CN–TiO ₂	HTT (180 °C for 4 h)	CO ₂ and H ₂ O	Simulated solar light	CO (NP)	NP	133
Core@shell <i>meso</i> -TiO ₂ @ZnIn ₂ S ₄ /Ti ₃ C ₂	Electrostatic self-assembly	CO ₂ and H ₂ O	Visible	CO (99.58 $\mu\text{mol g}^{-1}$ for 5 h) CH ₄ (8.97 $\mu\text{mol g}^{-1}$ for 5 h)	NP	134
Ti ₃ C ₂ QDs/Cu ₂ O NWs/Cu mesh	Dip coating and calcination under Ar (200 °C for 1 h)	CO ₂ and H ₂ O	Simulated solar light	CO (30.5 $\mu\text{mol g}^{-1}$ for 3 h) CH ₄ (34 $\mu\text{mol g}^{-1}$ for 3 h)	NP	135
Ti ₃ C ₂ sheets/Cu ₂ O NWs/Cu	Dip coating and calcination under Ar (200 °C for 1 h)	CO ₂ and H ₂ O	Simulated solar light	CH ₃ OH (78.50)	NP	135
Ti ₃ C ₂ /Ti ₃ C ₂ heterojunction composites	HTT (90 °C for 5 h)	CO ₂ and H ₂ O	Simulated solar light	CH ₃ OH (36.51)	NP	136
Cu ₂ O/Ti ₃ C ₂ heterojunction composites	HTT (120 °C for 24 h)	CO ₂	Simulated solar light	CO (17.55) CH ₄ (0.96)	NP	137
2D/2D Ti ₃ C ₂ /Bi ₂ WO ₆ nanosheets	HTT (120 °C for 24 h)	NaHCO ₃ and H ₂ SO ₄	Simulated solar light	CH ₄ (1.78) CH ₃ OH (0.44)	NP	138
Alkalinized Ti ₃ C ₂ /ZnO	KOH treatment followed by electrostatic self-assembly	CO ₂ and H ₂ O	Simulated solar light	CO (30.30) CH ₄ (20.33)	0.32 0.46	139
2D/2D Ti ₃ C ₂ /g-C ₃ N ₄ nanosheets	Calcination under N ₂ (550 °C for 2 h)	NaHCO ₃ and H ₂ SO ₄	Visible	CO (5.19) CH ₄ (0.044)	NP	140
2D/2D TiO ₂ /C ₃ N ₄ /Ti ₃ C ₂	Electrostatic self-assembly	NaHCO ₃ and H ₂ SO ₄	Simulated solar light	CO (4.39) CH ₄ (1.20)	NP	141
Alkalinized Ti ₃ C ₂ /g-C ₃ N ₄	Wet chemical mixing	CO ₂ and H ₂ O	Visible	CO (11.21 $\mu\text{mol g}^{-1}$ for 5 h) CH ₄ (0.203 $\mu\text{mol g}^{-1}$ for 5 h)	NP	142
Few layer Ti ₃ C ₂ /boron-doped g-C ₃ N ₄	Sonication and freeze drying	CO ₂ and H ₂ O	Visible	C ₂ H ₄ ; CH ₃ CHO CO (14.4 $\mu\text{mol g}^{-1}$ for 5 h) CH ₄ (0.8 $\mu\text{mol g}^{-1}$ for 5 h)	NP	143
g-C ₃ N ₄ /BiOIO ₃ /Ti ₃ C ₂ ternary composites	Electrostatic self-assembly	CO ₂ and H ₂ O	Visible	H ₂ (3.35 $\mu\text{mol g}^{-1}$ for 5 h) CO (5.88)	NP	144
3D hierarchical Co–Co LDH/Ti ₃ C ₂ T _x nanosheets	HTT (120 °C for 1 h)	MeCN, H ₂ O, TEA, [Ru(bpy) ₃]Cl ₂ and CO ₂	Visible	CH ₄ (1.55) CO (12.500)	0.92	145
3D hierarchical NiAl LDH/Ti ₃ C ₂	HTT (120 °C for 24 h)	CO ₂ and H ₂ O	Simulated solar light	H ₂ (NP) CO (11.82)	NP	146
0D/2D FAPbBr ₃ /Ti ₃ C ₂	Wet chemical synthesis	CO ₂ and H ₂ O	Simulated solar light	CH ₄ (1.02) CO (283.41)	NP	147
2D/2D FAPbBr ₃ /Ti ₃ C ₂	Wet chemical synthesis	CO ₂ and H ₂ O	Simulated solar light	CH ₄ (17.67) H ₂ (4.51)	NP	148
Cs ₂ AgBiBr ₆ nanocrystals/Ti ₃ C ₂ T _x nanosheets	Ultrasonication	CO ₂ and H ₂ O	Visible	CO (93.82) H ₂ (~0.9) CO (11.1 $\mu\text{mol g}^{-1}$) CH ₄ (1.3 $\mu\text{mol g}^{-1}$) H ₂ (8.9 $\mu\text{mol g}^{-1}$)	0.083	148

^a ~ Data taken from a graphical representation of the corresponding reference; NP: value not provided; MeCN: acetonitrile; TEA: triethanolamine.



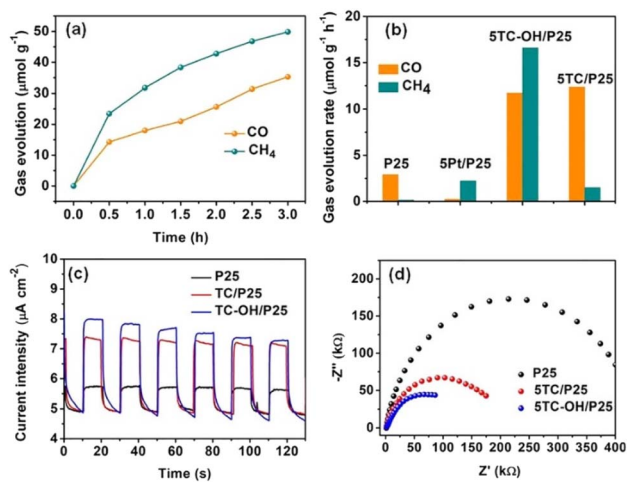


Fig. 10 (a) Gas evolution (CO and CH₄) over 5TC-OH/P25 as a function of irradiation time under a 300 W Xe lamp. (b) Evolution rates of CO and CH₄ over P25, 5Pt/P25, 5TC/P25, and 5TC-OH/P25 under irradiation of a 300 W Xe lamp. (c) Transient photocurrent responses and (d) EIS Nyquist plots of P25, 5TC/P25, and 5TC-OH/P25. Reproduced with permission from ref. 131, Copyright 2018 European Chemical Societies Publishing.

amounts of Ti₃C₂T_x, synthesized under hydrothermal conditions were studied for the photoconversion of CO₂ to CO in the absence of a hydrogen source.¹³⁶ Addition of 10 mg of Ti₃C₂T_x to the initial precursor solution for Cu₂O synthesis, exhibited 3 times higher CO production rates compared to that of pristine Cu₂O. After three cycles, the amount of CO produced remains 85.8% in comparison with the first cycle. The slight decrease was attributed to the loss of the sample during recovery. Surprisingly, the authors noticed the formation of minor amounts of CH₄. However, since the reaction took place under a pure CO₂ atmosphere, the detection of CH₄ clearly indicates the presence of impurities. Here also, the performance enhancement was attributed to the separation and concentration of electrons on Ti₃C₂, facilitating CO₂ conversion.¹³⁶

The 2D/2D heterojunctions were found to be very active in photocatalytic transformation of CO₂. Therefore, Cao *et al.*¹³⁷ fabricated a 2D/2D heterojunction comprising Ti₃C₂T_x/Bi₂WO₆ nanosheets under hydrothermal conditions. H₂SO₄ and NaHCO₃ were reacted for the *in situ* production of CO₂ which under simulated solar light irradiation produced CH₄ as the major product along with CH₃OH. The results indicate a 4.6 times increment in CH₄ and CH₃OH yields compared to that of pristine Bi₂WO₆ ultrathin nanosheets. The large interfacial surface area resulting from the 2D/2D heterojunction and lower charge transport distance ensured efficient electron transfer from Bi₂WO₆ to Ti₃C₂. This effect coupled with enhanced CO₂ adsorption capacity resulted in the observed performance of this study.¹³⁷

In the same line, Yang *et al.*¹³⁹ synthesized a series of 2D/2D Ti₃C₂/g-C₃N₄ ultra-thin heterojunctions by performing calcination of varying amounts of Ti₃C₂T_x and urea. Here, apart from being the precursor for C₃N₄, the presence of urea facilitated the exfoliation of MXene sheets. Photocatalytic conversion of *in situ*

generated CO₂ was carried out under visible light irradiation. In the absence of Ti₃C₂T_x, CO was the major product observed beside small traces of CH₄. The optimized composite material (with 10 mg of Ti₃C₂) added to the urea precursor, exhibited better yields of CO and CH₄ with 8.1 times higher CO₂ conversion, compared to pure C₃N₄. The performance, structure and morphology of the photocatalyst remained reasonably unchanged during five cycles. The detrimental effect of excess Ti₃C₂T_x was observed in this study also, as previously reported by Low *et al.*¹³⁰ The spatial separation of charge carriers facilitated by the formation of an ultra-thin heterojunction, along with an improved CO₂ adsorption and activation was responsible for the improvement of performance.¹³⁹

Integration of Ti₃C₂ quantum dots into a TiO₂/C₃N₄ core@shell structure resulted in a 2D/2D/0D dual heterojunction photocatalyst. This material exhibited superior photoconversion of *in situ* generated CO₂ into CO and CH₄ along with lower amounts of methanol and ethanol. Product yields remained consistent during four cycles of photocatalytic reactions. The formation of dual heterojunctions was found to be responsible for an improved separation and utilization of charge carriers, resulting in the enhanced performance.¹⁴⁰

The effect of alkalized Ti₃C₂ was also observed by coupling it with g-C₃N₄. The composite prepared by mixing 5 wt% alkalized Ti₃C₂ with g-C₃N₄ exhibited better CO evolution rates during photocatalytic conversion of a CO₂ saturated H₂O medium under visible light irradiation, compared to pure g-C₃N₄. A slight decrease in performance was observed during the first three cycles. However, after washing the catalyst, the activity increased and remained unchanged during the fourth and fifth cycles. A large Fermi level difference between the components contributed towards the enhanced separation of photogenerated charge carriers.¹⁴¹

Wang *et al.*¹⁴² synthesized a few-layer Ti₃C₂/B-doped C₃N₄ composite material by adding different volumes of an aqueous solution of Ti₃C₂T_x (1 mg mL⁻¹) to a fixed amount of B-doped g-C₃N₄ (0.4 g). Because of the intimate interfacial contact, the optimal (12 mL) MXene suspension added to the composite exhibited 3.2 times and 8.9 times higher CO and CH₄ yields, respectively, compared to pristine g-C₃N₄ under visible light irradiation. A slight decrease (~14%) in CH₄ and CO production was observed during the first three cycles which was then increased after washing with water.¹⁴²

A BiOIO₃/g-C₃N₄ Z-scheme heterojunction modified with Ti₃C₂ nanosheets was prepared by Hong *et al.*¹⁴³ following an electrostatic self-assembly method. Photocatalytic CO₂ conversion in an aqueous medium under visible light irradiation indicated a 6.6 times higher CO production rate for heterojunction photocatalysts with 4 wt% Ti₃C₂, in comparison with pure g-C₃N₄. No significant reduction in CH₄ and CO yields was observed during three cycles.¹⁴³

Chen *et al.*¹⁴⁴ synthesized a 3D hierarchical Co-Co layered double hydroxide/Ti₃C₂T_x nanoarray under HTT. These materials exhibited a significant enhancement in CO evolution rates, with an apparent quantum efficiency of 0.92%. The high electron conductivity of Ti₃C₂T_x, coupled with the nanoarray architecture synergistically contributed towards the



performance enhancement. For these materials, the product yield remained 90% of its initial value after five cycles.¹⁴⁴

Shi *et al.*¹⁴⁵ synthesized a 3D hierarchical NiAl layered double hydroxide/Ti₃C₂ nanocomposite under HTT by adding different volumes of an aqueous solution of Ti₃C₂ (15 mg mL⁻¹) to the NiAl-LDH precursor solution. Photoconversion of CO₂ was carried out using H₂O as the hydrogen source, producing CO with 92% selectivity, along with CH₄. Up to 3 times enhancement of CO production was observed and the 2D/2D interface interaction was found to promote charge separation and to facilitate CO₂ capture and activation. The CO yield remained constant for four cycles.¹⁴⁵

Organic–inorganic hybrid perovskites are another interesting category of materials coupled with MXenes for photocatalytic applications. Que *et al.*¹⁴⁶ anchored FAPbBr₃ QDs on Ti₃C₂ nanosheets resulting in a composite 0D/2D heterojunction photocatalyst for CO₂ reduction. The amount of Ti₃C₂ in the composite was varied between 0 and 1 mg. Under simulated solar light irradiation, CO was formed as the major product along with minor quantities of CH₄ and H₂ as shown in Fig. 11. In the presence of 0.2 mg of Ti₃C₂, CO yields were found to be 1.84 times higher compared that of to pure FAPbBr₃. No significant changes in product yields were observed during three reaction cycles. Ti₃C₂ QDs served as the electron acceptor as well as provided more active sites.¹⁴⁶

In another study, FAPbBr₃ nanoplates were combined with Ti₃C₂ by Que *et al.*¹⁴⁷ resulting in a 2D/2D Schottky heterojunction. A 1.5-fold enhancement in average CO production was observed for the composite, compared to pure halide perovskite.¹⁴⁷ Zhang *et al.*¹⁴⁸ assembled whole inorganic halide double perovskite Cs₂AgBiBr₆ nanocrystals (NCs) on Ti₃C₂T_x nanosheets. Superior product yields with 71% selectivity to CO, were obtained for the heterostructures, compared to the individual components. The authors used the electron consumption yield to determine the stability of the material. It remained almost unchanged after 3 cycles. Structural or morphological changes were also not observed during the reaction. The presence of Ti₃C₂T_x reduced the large exciton binding energy of Cs₂AgBiBr₆ and promoted the formation of free charge carriers with prolonged life times. Charge carrier recombination was therefore suppressed and this together with enhanced CO₂ adsorption, resulted in better performance.¹⁴⁸

3.5. Photocatalytic applications for N₂ photofixation

In addition to the well-established photocatalytic applications presented above, the MXenes are used as a cocatalyst in photocatalytic systems for the N₂ reduction reaction (NRR) to ammonia. In contrast to modern Haber-Bosch technology, which directly utilizes fossil fuels through intensive energy processes at high pressure and temperature and with high CO₂ emissions,^{149,150} the NRR is regarded as an environmentally friendly, sustainable and green NH₃ synthesis that utilises N₂ and water molecules under mild conditions.¹⁵¹ Obtaining high reaction efficiencies is primarily constrained by electron–hole recombination and limited interfacial charge transfer, but it is also difficult to activate and break the strong and stable NN

bond.¹⁵² The effectiveness of the NRR can be increased by using MXenes as a cocatalyst to create Schottky junctions with semiconductor materials. Table 3 highlights a few examples of MXene-based photocatalysts that have been applied recently for the NRR.

The 0D/2D AgInS₂/Ti₃C₂ Z-scheme heterostructures constructed by HTT, with various mass ratios of AgInS₂, showed a notable improvement in photocatalytic activity for the NRR, according to Qin *et al.*¹⁵³ For the mass ratio of 30% AgInS₂ in N₂ saturated solution and under visible light, the greatest ammonia yield rate was 38.8 μmol g⁻¹ h⁻¹, with an AQY = 0.07% at 420 nm. The utilization of Ti₃C₂ MXenes as a cocatalyst and Z-scheme heterojunction, allowed the effective charge separation and migration of photogenerated electrons and holes, which are responsible for the photocatalytic activity for the NRR.

According to Chang *et al.*,¹⁵⁴ the usage of partly reduced Ti₃C₂ (r-Ti₃C₂) with Au nanospheres interlaminated in a layered r-Ti₃C₂ structure helped to expose several low-valence Ti sites that act as active sites. Injection of photogenerated hot electrons from Au nanospheres under illumination reduces N₂ molecules, which are then captured and activated at the active site. In N₂ saturated water, the NRR of r-Ti₃C₂/Au generated NH₄⁺ at a rate of 12.4 μmol g⁻¹ h⁻¹ under visible light, which is 5.9 and 10.3 times higher than those of Ti₃C₂/Au and r-Ti₃C₂, respectively, while under white light the rate was 22.6 μmol g⁻¹ h⁻¹, which is 5.8 and 10.2 times higher than those of Ti₃C₂/Au and r-Ti₃C₂. By measuring the quantity of ammonia generated in pure water under monochromatic light irradiation, the apparent quantum efficiencies (AQE) of r-Ti₃C₂/Au were found to reach 0.697% at 520 nm. The NH₃ production rate increased to 21.26 μmol g⁻¹ h⁻¹ when the lighting source was changed to simulated sunlight irradiation (by using an AM 1.5G filter).

The material exhibits high stability, with ammonia generation remaining close to 95%, according to the results of five successive stability test cycles. This may be due to the electrons that are constantly produced and transferred from the Au nanospheres to Ti₃C₂ under illumination, which can prevent oxidation of MXenes.

Hao *et al.*¹⁵⁵ demonstrated outstanding activity for the NRR in water at room temperature using a RuO₂-loaded TiO₂-MXene

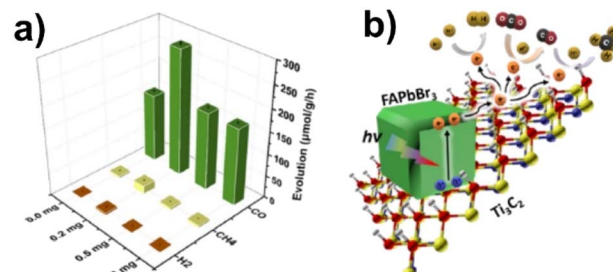


Fig. 11 (a) Photocatalytic CO₂ reduction performances of FAPbBr₃/x-Ti₃C₂ (x = 0.0, 0.2, 0.5, and 1.0) in deionized water under AM 1.5 G (150 mW cm⁻²). (b) Schottky heterojunction photocatalytic system for CO₂ reduction in FAPbBr₃/x-Ti₃C₂. Reproduced with permission from ref. 146, Copyright 2021 American Chemical Society.



Table 3 Compilation of synthetic and photocatalytic aspects of MXene-based materials in N₂ photofixation^a

Photocatalyst	Preparation methods	Reaction conditions	Light source	Sacrificial reagent	NH ₃ production [$\mu\text{mol g}^{-1} \text{h}^{-1}$]	AQY [%]	Ref.
0D/2D AgInS ₂ /Ti ₃ C ₂	HTT (150 °C for 5 h)	20 mg of sample dispersed in 100 mL 20% methanol solution, RT	300 W xenon lamp (>400 nm)	Methanol	38.8	0.07 (420 nm)	153
r-Ti ₃ C ₂ /Au	Solvent-driven	80 mg of the photocatalyst and 50 mL of water, RT 200 mg of the photocatalyst and 50 mL of water, RT 50 mg of photocatalyst with 100 mL distilled water	300 W xenon lamp Simulated solar light	None	22.6 ^b 12.4 ^c 21.26	0.697 (520 nm)	154
RuO ₂ @TiO ₂ -MXene	HTT (150 °C for 10 h)	30 mg was dispersed in 100 mL of ultrapure water	300 W xenon lamp, full spectrum (100 mW cm ⁻²) 300 W xenon lamp (UV-vis light)	None	4.37	NP	155
MXene/TiO ₂ /Co-0.5%	Two-step calcination under N ₂ (400 °C for 1 h) HTT (200 °C for 5 h)	30 mg was dispersed in 100 mL of ultrapure water 20 mg photocatalyst dispersed in 100 mL aqueous solution containing 20 vol% methanol	300 W xenon lamp	None	106.7	NP	156
Cds@Ti ₃ C ₂	HTT (200 °C for 5 h)	20 mg photocatalyst dispersed in 100 mL aqueous solution containing 20 vol% methanol	300 W xenon lamp	Methanol	293.06 $\mu\text{mol L}^{-1} \text{h}^{-1}$	7.88	113
Nb ₂ O ₅ /C/Nb ₂ C/g ⁻¹ -C ₃ N ₄	Calcination under CO ₂ (850 °C) and co-sintering (550 °C for 2 hours)	50 mg of photocatalyst in 50 mL of double distilled water with 20 vol% methanol	Visible light ($\lambda > 420 \text{ nm}$)	Methanol	365	NP	157
TiO ₂ @C g ⁻¹ -C ₃ N ₄ :10%	One-step calcination in air (550 °C for 2 hours)	50 mg of photocatalyst suspended in 100 mL of DD water containing 20 vol% methanol	300 W xenon lamp	Methanol	250.6	0.14 (420 nm)	158
OV-C/TiO ₂ -600	One-step calcination in air (600 °C for 2 hours)	50 mg of catalyst dispersed in 50 mL of ultrapure water, RT	300 W xenon lamp (200–800 nm)	H ₂ O Methanol	41 81	0.04 (400 nm)	159
Ti ₃ C ₂ T _x /TiO ₂ -400	Calcination in air (400 °C for 15 min)	10 mg of photocatalysts and 20 mL of deionized water, RT	Xenon lamp (320–780 nm), full spectrum (250 mW cm ⁻²)	None	422	NP	160
W/Ti ₃ C ₂ T _x -U	Ball milling	20 mL of ultrapure water, 10 mg of purified photocatalysts, RT	Xenon lamp, full-spectrum (250 mW cm ⁻²)	None	227.5	0.05–0.15 in UV-vis and NIR	161
C ₃ N ₄ /r-Ti ₃ C ₂ QDs-2	Electrostatic assembling	20 mg photocatalyst and 50 mL of 20 vol% methanol, RT	300 W xenon lamp	Methanol	328.9 ^b 197 ^c	0.92 (380 nm)	162

^a NP: value not provided. ^b White light (300–780 nm). ^c Visible light (420–780 nm).



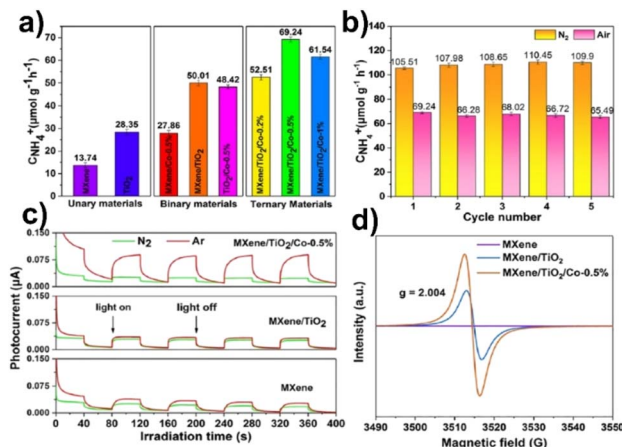


Fig. 12 (a) NH₄⁺ production rate in ambient air; (b) the stability of N₂ photofixation activity of MXene/TiO₂/Co-0.5% under N₂ and an air atmosphere; (c) photocurrent spectra of MXene, MXene/TiO₂ and MXene/TiO₂/Co-0.5%; (d) EPR spectra. Reproduced with permission from ref. 163, Copyright 2015 American Chemical Society.

hybrid nanostructure with varied RuO₂ ratios. After 450 minutes of Xe-lamp irradiation, the as-prepared RuO₂@TiO₂-MXene exhibits good N₂ reduction performance, reaching a production rate of 4.37 μmol g⁻¹ h⁻¹. The results show that RuO₂@TiO₂-MXene is really a viable material for the NRR. Since TiO₂ creates electrons and holes, MXenes offer efficient electron transport due to metallic conductivity, and RuO₂ presents activation sites, and combining the three components in the hybrid structure has a significant impact on the material's photofixation performance.

In a two-step calcination procedure, Gao *et al.*¹⁵⁶ created MXene/TiO₂/Co composite materials for the NRR. The production rate of MXene/TiO₂/Co-0.5% increased 1.4 times by the addition of Co (50 μmol g⁻¹ h⁻¹) compared to that of MXene/TiO₂, and 2.4 times compared with that of TiO₂ (69 μmol g⁻¹ h⁻¹). It is significant to note that the ammonia concentration reached 107 μmol g⁻¹ h⁻¹ under UV-vis light. More than this, the photocatalysts exhibit structural stability throughout the reaction as the XRD diffractograms after five cycles of the NRR show no change at all before and after 10 hours of irradiation. Electron paramagnetic resonance (EPR) spectroscopy was used to examine the reaction mechanism and the results suggested that MXene/TiO₂/Co-0.5% had a higher oxygen vacancy concentration which is beneficial for photocatalytic nitrogen fixation,¹⁶³ as shown in Fig. 12.

Sun *et al.*¹¹³ investigated the effects of Ti₃C₂ as a co-catalyst generated by HTT for the NRR, as well as the photocatalytic performance of CdS@Ti₃C₂ composites. Pure CdS NRs (nanorods) had a low rate of photocatalytic NH₄⁺ formation (156.62 μmol L⁻¹ h⁻¹), while Ti₃C₂ does not produce any ammonia. To create the CdS@Ti₃C₂ composites, several amounts of Ti₃C₂ ranging from 15 to 100 mg, were tested. Of these, in the presence of a scavenger, CdS@Ti₃C₂-15 with 15 mg of MXenes has shown the best NRR activity at a rate of 293.06 μmol L⁻¹ h⁻¹, which is 1.9 and 1.4 times higher than that of pure CdS NRs and

CdS@Pt, respectively, with an AQY = 7.88%. As a result, Ti₃C₂ MXenes may be able to replace and outperform Pt in terms of enhancing CdS's capacity for the NRR. Photocatalytic nitrogen fixation on a CdS Ti₃C₂-15 composite was conducted in the absence of a sacrificial agent at a nitrogen fixation rate of 27 μmol L⁻¹ h⁻¹. The stability test revealed that the production rate is 99.35% of the initial rate after three cycles. Due to their affordability and stability, these composites are expected to play a significant role in the production of ammonia in the future.

Jiang *et al.*¹⁵⁷ reported a 2D MXene-derived Nb₂O₅/C/Nb₂C/g-C₃N₄ heterojunction which was created by uniformly growing Nb₂O₅ on Nb₂C and then forming g-C₃N₄ nanosheets *in situ* on Nb₂O₅/C/Nb₂C. With methanol acting as a scavenger and an optimum Nb₂O₅/C/Nb₂C/g-C₃N₄ ratio of 1:1, the heterojunctions exhibited a 9.1 times higher nitrogen reduction rate (365 μmol g⁻¹ h⁻¹) in water under visible light, compared with an MXene derived Nb₂O₅/g-C₃N₄ composite. Additionally, a pH adjustment from 3 to 9 would enhance the efficiency of the NRR from 180 to 927 μmol L⁻¹ h⁻¹ by changing the energy barrier of proton reduction. The improved nitrogen reduction photocatalytic performance may be due to both the Schottky junction produced at the Nb₂O₅/Nb₂C interface as well as the increased electron and hole separation due to the close contact between the conductive Nb₂C and the Nb₂O₅. The NRR efficiency decreased as the pH of the catalytic system is higher than 9, due to extra negative charges created on the photocatalyst surface that prevent the reaction substrate from effectively making contact with the catalytically active site¹⁶⁴ (Fig. 13). Nb₂O₅/C/Nb₂C/g⁻¹-C₃N₄ heterojunctions are stable materials, with no noticeable NH₃ production rate loss observed after 5 cycles.

Through one-step calcination in air of a mixture of Ti₃C₂ and melamine with various ratios of TiO₂@C to g-C₃N₄, Liu *et al.*¹⁵⁸ fabricated a TiO₂@C/g-C₃N₄ heterojunction for the NRR. This approach produces a large number of Ti³⁺ species which act as active sites. The composite material involves the formation of TiO₂ by thermal treatment of Ti₃C₂T_x MXenes, which is further tightly wrapped by *in situ* formed g-C₃N₄ nanosheets. The remarkable performance of the TiO₂@C/g-C₃N₄ heterojunction for photocatalytic nitrogen reduction to ammonia is due to its extensive surface defects, high electron-donating ability, improved light absorption zone, efficient charge transport and reduced recombination, and also strong nitrogen activation ability. The TiO₂@C/g-C₃N₄:10% sample had the highest photocatalytic activity of all the samples, with an ammonia production rate of 250.6 μmol g⁻¹ h⁻¹, with an AQE of 0.14% (420 nm) under visible light irradiation. The performance of TiO₂@C/g-C₃N₄:10% is found to be 18 and 10 times higher than that of g-C₃N₄ and TiO₂@C, respectively. A recycling NH₃ test on TiO₂@C/g-C₃N₄ revealed no detectable decrease in the NH₃ production rate following five cycles in six hours, demonstrating the good stability of the sample.

Using a one-step calcination method at varied temperatures, Qian *et al.*¹⁵⁹ used Ti₃C₂ as a precursor to produce an oxygen vacancy (OV)-rich C/TiO₂ (OV-C/TiO₂) material to convert N₂ into NH₃ under sunlight irradiation. Among the samples, OV-C/TiO₂-600 produced NH₃ at a rate of 41 μmol g⁻¹ h⁻¹ using water as a proton source (AQY of 0.04% for 400 nm light and 0.01% for



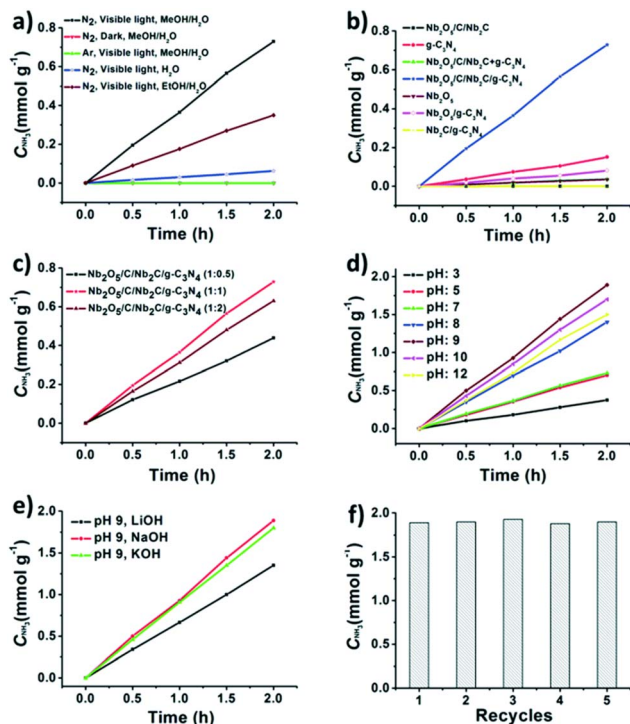


Fig. 13 (a) Control experiments on photocatalytic NH₃ production with the MXene-derived Nb₂O₅/C/Nb₂C/g-C₃N₄ heterojunction under different conditions. (b) Photocatalytic activity of the Nb₂O₅/C/Nb₂C, g-C₃N₄, Nb₂O₅/C/Nb₂C + g-C₃N₄, Nb₂O₅/C/Nb₂C/g-C₃N₄, Nb₂O₅, Nb₂O₅/g-C₃N₄ and Nb₂C/g-C₃N₄ for NH₃ production under visible light irradiation ($\lambda > 420$ nm). (c) Photocatalytic NH₃ production activity of Nb₂O₅/C/Nb₂C/g-C₃N₄ with the ratio variation of Nb₂O₅/C/Nb₂C: melamine from 0.5 to 1 and 2 during catalyst preparation; (d) photocatalytic NH₃ production activity of Nb₂O₅/C/Nb₂C/g-C₃N₄ with different pH values adjusted by using HCl or NaOH solution. Reproduced with permission from ref. 157, Copyright 2020 The Royal Society of Chemistry.

of 420 nm light, respectively) which is 26.5 times higher than that of TiO₂ Degussa P25. When using methanol as a proton source the NH₃ rate increased to 84 $\mu\text{mol g}^{-1} \text{h}^{-1}$. The addition of oxygen vacancies on the TiO₂ surface and carbon doping increased electron-hole pair formation and decreased electron-hole pair recombination. Moreover, there was an increase in the photocarrier lifetime, as well as chemisorption and activation of N₂ molecules. The NH₃ production yields essentially stayed constant during four photocatalytic cycling trials in 2 h, showing the OV-C/TiO₂-600 sample's good stability.

Hou *et al.*¹⁶⁰ developed plasmonic Ti₃C₂T_x and TiO₂ hybrid structures (Ti₃C₂T_x/TiO₂) at various temperatures, with Ti₃C₂T_x/TiO₂-400 exhibiting higher activity for the NRR. At room temperature, the Ti₃C₂T_x/TiO₂-400 hybrid structure yielded 422 $\mu\text{mol g}^{-1} \text{h}^{-1}$ of NH₃ without the use of any sacrificial agents. After being exposed to monochromatic light at 630 nm, Ti₃C₂T_x/TiO₂-400 generated 50 $\mu\text{mol g}^{-1} \text{h}^{-1}$ of NH₃, with an AQE of 0.05%. Once monochromatic light was changed to 740 nm light, the NH₃ generation rate for Ti₃C₂T_x/TiO₂-400 improved to 82 $\mu\text{mol g}^{-1} \text{h}^{-1}$, while the AQE became 0.07%. Additionally, the stability test of Ti₃C₂T_x/TiO₂-400 shows that more than 90% of

the initial reaction activity was conserved in ten consecutive rounds of reaction, demonstrating the high stability of Ti₃C₂T_x/TiO₂-400. The activity of Ti₃C₂T_x/TiO₂-400 hybrid structures was attributed to the coexistence of Ti₃C₂T_x and defective TiO₂, which considerably improved charge carrier separation efficiency. The plasmonic Ti₃C₂T_x phase may collect visible and near-infrared light to generate hot electrons that are transported to the TiO₂ conduction band and then trapped by oxygen vacancies.

Under full-spectrum irradiation at room temperature without the use of sacrificial materials, Qi *et al.*¹⁶¹ demonstrated a remarkable ammonia production rate (227.5 $\mu\text{mol g}^{-1} \text{h}^{-1}$) utilizing Ti₃C₂T_x doped with a W atom. Further research on the catalytic performance was conducted using monochromatic light of various wavelengths. At 630 nm, the production rate was 43.6 $\mu\text{mol g}^{-1} \text{h}^{-1}$, but it dropped to 28 $\mu\text{mol g}^{-1} \text{h}^{-1}$ when the light source was centered at 740 nm. In particular, the AQEs of W/Ti₃C₂T_x-U were 0.05–0.15% in UV-vis and NIR wavelengths. In addition, W/Ti₃C₂T_x-U are stable materials preserving >90% activity after ten subsequent rounds of reaction. Doping with W atoms enhances the physical and chemical adsorption of N₂, and they also serve as the active sites for reducing reaction energy during the NRR.

According to Chang *et al.*,¹⁶² mesoporous hollow g-C₃N₄ spheres were improved with partly reduced Ti₃C₂ quantum dots (r-Ti₃C₂ QDs) on the surface for effective NRR. This resulted in the formation of a C₃N₄/r-Ti₃C₂ QD Schottky junction *via* electrostatic self-assembly. The C₃N₄/r-Ti₃C₂ QDs' nitrogen photo-fixation activity was better when exposed to white light than it was when exposed to visible light since the photocatalysts had a limited capacity to absorb visible light. The ammonia

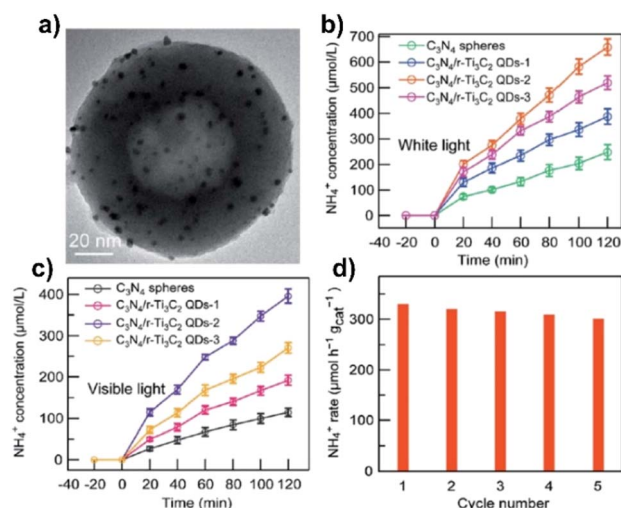


Fig. 14 (a) TEM image of C₃N₄/r-Ti₃C₂ QDs-2; (b and c) N₂ photofixation over the different as-obtained photocatalysts. Time courses of the ammonia concentrations measured under white light illumination (light intensity: 300 mW cm⁻² and $\lambda = 300$ –780 nm) and visible light illumination (light intensity: 300 mW cm⁻² and $\lambda = 420$ –780 nm); (d) stability tests of C₃N₄/r-Ti₃C₂ QDs-2 for N₂ photofixation under white light. Reproduced with permission from ref. 162, Copyright 2022 The Royal Society of Chemistry.



production rate of $C_3N_4/r-Ti_3C_2$ QDs reduced from $328.9 \mu\text{mol g}^{-1} \text{h}^{-1}$ to $197 \mu\text{mol g}^{-1} \text{h}^{-1}$ when the light source was switched from white to visible light (Fig. 14), which, under the same circumstances, are 1.6 and 1.9 times as high as that of C_3N_4 , respectively. The AQE value of $C_3N_4/r-Ti_3C_2$ QDs is 0.92% at 380 nm. When C_3N_4 is exposed to light, photoinduced electrons and holes are produced, and $r-Ti_3C_2$ QDs enable effective migration and separation for these particles. Numerous OV-defect state sites and Ti^{3+} sites found in $r-Ti_3C_2$ QDs serve as active sites for collecting and activating N_2 molecules. Quickly moving from C_3N_4 to $r-Ti_3C_2$ QDs *via* the close contact interface, photogenerated electrons from C_3N_4 may next be reduced to NH_3 by trapping and accumulating photoexcited electrons from $g-C_3N_4$.

4. Metal oxides

4.1. Synthesis and structure evolution

The most facile strategy to tailor the physicochemical properties of MOs is by varying their composition. In binary oxides such as TiO_2 , ZnO , Fe_2O_3 , *etc.*, a small quantity of an additional metallic component can be introduced into the lattice, while maintaining the overall crystal structure.^{165–167} This approach is more straightforward in mixed oxides such as perovskites, brownmillerites, *etc.*, since such materials include more than one metal in their pristine form.

However, more complex compositions with fascinating properties can be made by doping with additional cationic components.^{168–171} In all these cases, excellent control of desired properties, including photocatalysis, can be achieved. On the other hand, photocatalytic performance also significantly relies on the nanoscale structural features and morphology of the materials involved. Since fabricating nanostructures depends on synthesis methodology, the choice of the synthesis method is important. The conventional approach for making MOs is the solid-state method where thermal energy drives the formation of the desired phase with a high degree of crystallinity. Here, agglomerated large grains are often produced with lower values of specific surface area. Mechanochemical approaches such as ball milling replaced thermal energy with mechanical energy, with limited extent of applicability and poor control over morphology and nanostructure. It is to be noted that mixed oxide compositions more or less require higher temperatures for the formation of the desired single-phase materials. Also, it is difficult to ensure a homogeneous mixing of the precursors in solid state and mechanochemical methods.^{168–171} Therefore, several soft chemical approaches were developed, incorporating a variety of precursors, templates and reaction media by following a number of complexation, precipitation, hydrothermal and templating routes.^{172–175} Wet chemical approaches such as the polyol method, sol-gel method, *etc.*, allowed the synthesis of semiconductor photocatalysts with a uniformly distributed particle size and morphology, at comparatively lower temperatures. These methods generally involve the presence of organic molecules such as citric acid, ethylene glycol, *etc.* The interactions between these organic components with their metallic counterparts along with the mechanism of their

cross linking and/or decomposition can exert better control on particle size and morphology.¹⁷²

One step synthesis of binary and ternary oxides can be achieved by carrying out the reaction in a sealed reaction vessel above the boiling point of the solvent. This method is designated as the hydrothermal method when water is used as the solvent or the solvothermal method when an organic solvent is used instead.^{173,176} A homogeneous solution of the precursor in an appropriate solvent is subjected to thermal treatment at a temperature higher than the boiling point of the solvent. This promotes an increase in pressure inside the reaction chamber. The combined effect of enhanced temperature and autogenous pressure will drive the formation of the desired product. Excellent control of morphology and nanostructure formation is observed in this method. Simple synthesis parameters such as reactant concentration, pH and temperature were found to significantly influence the formation of the desired phase and morphology in most of the cases. Additional control of microstructure and porosity can be achieved by introducing appropriate amounts of organic templates during synthesis. Even though an additional calcination step is not necessary in most of the cases, high temperature treatments are always required to remove the organic structure directing agents and for the structure stabilization of the materials for applications involving enhanced temperatures.¹⁷⁷ To date, a variety of binary and ternary oxide compositions with varied nanostructures and morphologies have been synthesized by following the hydrothermal method.^{178–180}

Followed by the successful synthesis of MCM-41 silica with ordered mesopore channels, templating methods are widely used to produce MOs with controlled porosity and textural parameters.¹⁸¹ The initial focus was on the use of surfactants as soft templates to direct the synthesis of porous oxides.¹⁸² Apart from ordered mesoporous silica and carbon materials, limited success was obtained for the soft template method for oxides.^{183–185} This was primarily due to the non-availability of precursors that undergo hydrolysis in a controlled manner with organic templates. Also, organic template removal is generally carried out at high temperatures. Since the crystallization temperature for most of the MO phases was higher than the decomposition temperature of the template, the quality of the resultant materials was compromised. On the other hand, the nanocasting method was also being developed in parallel, for which an inert porous oxide was used as the hard template.^{182–187} This method was initially used for the synthesis of ordered mesoporous carbon and was then extended to oxides, utilizing ordered mesoporous silica or carbon as hard templates. A wide variety of binary oxides with extremely high values of surface area and well-ordered pore structure were obtained by this method. Nanocasting was found to be highly versatile and was then used to synthesize complex ternary oxides and non-oxide compositions such as sulfides.^{188,189} Unfortunately, this multi-step method was time consuming, costly and non-scalable.

Uniform integration of oxide cocatalysts with semiconductors is necessary to develop highly efficient photocatalysts. Several studies reported single step calcination



treatments under oxidizing or inert atmospheres to combine oxide cocatalysts with photocatalysts including photocatalysts such as TiO₂, g-C₃N₄, *etc.*^{190–192} In addition, other approaches including simple impregnation of the cocatalyst species, HTT, photodeposition, *etc.*, are also followed to develop cocatalyst-photocatalyst combinations.^{193–195}

4.2. Photocatalytic applications for water splitting

MOs are considered as low cost and easily available cocatalysts for photocatalytic applications including H₂ evolution by splitting water. Fundamentally, the photogenerated electrons from the semiconductor photocatalyst move towards the MO, acting as a cocatalyst, through the interface, resulting in electron-hole separation. Table 4 summarizes several examples of such cocatalyst/photocatalyst systems used in the most recent years for hydrogen production through water splitting. The synthetic approaches to combine different semiconductor photocatalysts and several reaction parameters are also presented.

Thus, Chen *et al.*¹⁹⁶ photodeposited Ni₂O₃ on the surface of hydrothermally synthesized CdS, and H₂ evolution was monitored under visible light irradiation. The presence of Ni₂O₃

resulted in the enhancement of the hydrogen evolution rate up to 41 times that of pure CdS and up to 1.9 times that of Pt loaded CdS. The authors attributed this improvement to the ability of Ni₂O₃ to act as electron traps, thereby promoting charge separation. Stability tests were performed for five cycles and the activity was found to decrease to 86.8% of the initial activity. The authors attributed this to the catalyst loss during filtration and washing.¹⁹⁶ To study the effect of the chemical environment of photodeposition, nickel oxides were loaded onto CdS under neutral as well as alkaline conditions. Interestingly, the materials synthesized under alkaline conditions resulted in a 117 times enhancement of the hydrogen production rate compared to that of pure CdS. It was proposed that the presence of OH⁻ ions contributed to charge separation by facilitating hole transfer towards the sacrificial agent, methanol.¹⁹³

Lin *et al.*³⁰ combined 1D Cd_{1-x}Zn_xS nanocrystals with oxygen enriched MoS₂ resulting in Cd_{1-x}Zn_xS@O-MoS₂ nanohybrids. NiO_x species were deposited by following an *in situ* photodeposition method. The optimized composition of these nanohybrids exhibited 25 times higher hydrogen evolution rates compared to Pt containing CdZnS. Under visible light

Table 4 Compilation of synthetic and photocatalytic aspects of MO-based materials in hydrogen production through the water splitting reaction^a

Cocatalyst/photocatalyst	Synthetic method	Hole scavenger	Light region	Optimum H ₂ production rate [μmol h ⁻¹ g ⁻¹]	AQY [%]	Ref.
NiO _x /CdS	<i>In situ</i> photodeposition	Methanol (30%)	Visible	445.6	NP	196
NiO _x /CdS	<i>In situ</i> photodeposition	Methanol (30%)	Visible	590.8	8.6 (400 nm)	193
NiO/Zn _{0.8} Cd _{0.2} S	<i>In situ</i> photodeposition	None	Visible	783.11	5.86 (350 nm)	197
NiO/TiO ₂ nanofibers	Electrospinning and calcination in air (500 °C for 3 h)	Methanol (25%)	Full spectrum	377	1.7 (365 nm)	200
Amorphous NiO/g-C ₃ N ₄	Calcination in air (100–500 °C for 3 h)	TEA (10%)	Visible	68.8	~0.12 (370 nm)	190
NiO/g-C ₃ N ₄	Impregnation, HTT (125 °C for 10 h) and calcination in air (200–400 °C for 1 h)	None	Visible	1.41	NP	198
NiO/TiO ₂	Impregnation and calcination in static air (450 °C for 3 h)	Methanol (5%)	Simulated solar light	~130	NP	199
NiO/TiO ₂	Impregnation and calcination in static air (450 °C for 3 h)	Methanol (5%)	UV	~70	NP	199
NiO/red phosphorous	Calcination in air (300 °C for 1 h)	Methanol (10%)	Visible	57.27	0.37 (420 nm)	201
Amorphous MoO _x /CdS nanorods	Sonication	Lactic acid (10%)	Simulated solar light	19 300	21.3 (420 nm)	202
Cu ₂ O/g-C ₃ N ₄	Calcination in air (520 °C for 4 h)	TEA (10%)	Visible	241.3	NP	203
CuO/TiO ₂ composite nanofibers	Electrospinning and calcination in air (500 °C for 2 h)	Methanol (10%)	Simulated solar light	1146.9	NP	204
CuO nanodots on TiO ₂ nanocrystals	Precipitation using ammonium hydroxide followed by calcination in air (200 °C for 1 h)	Methanol (10%)	UV-visible	~16 000	NP	191
CuO@NiO core@shell nanoparticles	Chemical reduction (using N ₂ H ₄ ·H ₂ O) followed by calcination under an inert atmosphere (400 °C for 2 h)	Glycerol (5%)	Solar light	26 100	NP	192
CuO quantum dots/TiO ₂ nanosheets	Impregnation followed by calcination in air (300 °C for 2 h)	Methanol (20%)	Full spectrum	18 390	26.93 (313 nm)	205
CoO/g-C ₃ N ₄	Calcination under Ar (400 °C for 4 h)	TEA (10%)	Visible	651.3	NP	206
CoO/C ₃ N ₄ nanotubes	Calcination under vacuum (400 °C for 4 h)	TEA (10%)	Visible	788.6	4.9 (420 nm)	207
Co ₃ O ₄ /C ₃ N ₄ nanotubes	Calcination in static air (400 °C for 4 h)	TEA (10%)	Visible	725.7	4.07 (420 nm)	207

^a ~ Data taken from a graphical representation of the corresponding reference; NP: value not provided; TEA = triethanolamine.



irradiation using $\text{Na}_2\text{S}/\text{Na}_2\text{SO}_3$ as a sacrificial agent, an AQY of 64.1% was obtained for these materials. Repeated reactions during five cycles indicated minor activity loss. Also, the structure and composition of the recovered material remained unchanged after the photocatalytic reaction.³⁰

Liu *et al.*¹⁹⁰ synthesized a series of amorphous NiO modified $\text{g-C}_3\text{N}_4$ under different temperature treatments. The heterojunctions formed at the interface between amorphous NiO and $\text{g-C}_3\text{N}_4$ created an inner electric field allowing the transfer of photogenerated electrons from C_3N_4 to NiO. Under visible light irradiation, NiO/ $\text{g-C}_3\text{N}_4$ exhibited 430 times higher hydrogen evolution rates compared to pristine $\text{g-C}_3\text{N}_4$ and was consistent during five cycles. Also, the structure and chemical state did not change after the reaction. The authors concluded that the presence of amorphous NiO enhanced the visible light response, promoted the separation and transfer of charge carriers and provided active sites for hydrogen evolution.¹⁹⁰

Ning *et al.*¹⁹⁷ synthesized a series of $\text{Zn}_{1-x}\text{Cd}_x\text{S}$ and an ultrathin layer of NiO was assembled over it by photodeposition. Under visible light irradiation and in the absence of a sacrificial agent, these materials showed significantly higher photocatalytic activity with an AQY of 0.66% for H_2 evolution, compared to Pt/ ZnCdS , as shown in Fig. 15. The NiO layer suppressed the photocorrosion of ZnCdS and the strong electronic coupling between the NiO layer and $\text{Zn}_{1-x}\text{Cd}_x\text{S}$ enabled efficient charge separation and transport.¹⁹⁷

Fu *et al.*¹⁹⁸ synthesized a series of NiO- $\text{g-C}_3\text{N}_4$ by following a successive hydrothermal and calcination procedure. Overall water splitting was achieved for these materials among which the best performance was observed for the materials calcined at 300 °C. In this case, a H_2 evolution rate of $1.41 \mu\text{mol g}^{-1} \text{h}^{-1}$ and an O_2 evolution rate of $0.71 \mu\text{mol g}^{-1} \text{h}^{-1}$ were obtained. Experiments carried out for seven cycles along with characterization performed using the recovered catalyst confirmed the stability of these materials. Photocatalytic efficiency decreased for the materials calcined at higher temperatures. Here also, the presence of NiO aided in visible light absorption, separated the charge carriers and provided active sites.¹⁹⁸ These results are in agreement with the report by Liu *et al.*¹⁹⁰

Banic *et al.*¹⁹⁹ synthesized a series of NiO/ TiO_2 materials with varying mass ratios, by wet impregnation. The photocatalytic water splitting reaction was performed in the presence of methanol under simulated solar irradiation and artificial UV irradiation. The materials with 7 wt% NiO enhanced hydrogen production rates up to 7.6 times compared to that of pure TiO_2 and 112.8 times compared to that of pure NiO. Under optimum conditions, a 1.8 times higher hydrogen production rate was achieved under simulated solar light compared to that under the influence of artificial UV radiation.¹⁹⁹

Since the preparation method is very important, Li *et al.*²⁰⁰ synthesized mesoporous NiO/ TiO_2 nanofibers by the electrospinning method. The presence of NiO inhibited the crystal growth of TiO_2 resulting in an increase in surface area, pore volume and pore size. Under simulated solar irradiation, an optimum NiO content (0.25 wt%) exhibited a 7 times higher H_2 evolution rate with an AQY of 1.7%, compared to pure TiO_2 .²⁰⁰ Chen *et al.*²⁰¹ synthesized a NiO/RP (red phosphorous)

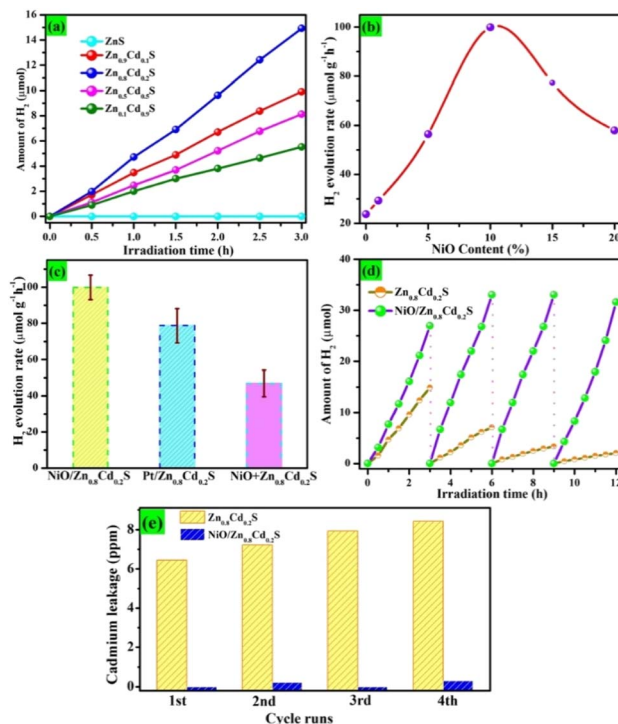


Fig. 15 (a) Photocatalytic H_2 evolution over $\text{Zn}_{1-x}\text{Cd}_x\text{S}$. (b) Rate of H_2 evolution over the x% NiO/ $\text{Zn}_{0.8}\text{Cd}_{0.2}\text{S}$ composite samples. (c) A comparison of the H_2 evolution rate over 10% NiO/ $\text{Zn}_{0.8}\text{Cd}_{0.2}\text{S}$, 2% Pt/ $\text{Zn}_{0.8}\text{Cd}_{0.2}\text{S}$, and NiO + $\text{Zn}_{0.8}\text{Cd}_{0.2}\text{S}$ (simple mechanical mixture of NiO and $\text{Zn}_{0.8}\text{Cd}_{0.2}\text{S}$). (d) Cycling runs for photocatalytic H_2 evolution in pure water with the $\text{Zn}_{0.8}\text{Cd}_{0.2}\text{S}$ and 10 wt% NiO/ $\text{Zn}_{0.8}\text{Cd}_{0.2}\text{S}$ composite samples under visible light irradiation. (e) Cd^{2+} leakage profiles under visible light irradiation. Reproduced with permission from ref. 197, Copyright 2019 European Chemical Societies Publishing.

composite by HTT followed by high temperature calcination. In this study, the 3 wt% NiO/RP composite exhibited the highest photocatalytic activity for hydrogen production which is 68.56 times higher than that of pure RP under visible light irradiation. The rates did not change significantly for 10 cycles and the physicochemical properties of the recovered materials were in agreement with those of the fresh ones. In the presence of light, electrons are excited towards the conduction band of RP leaving behind holes in the valence band, which simultaneously combines with the electrons from NiO. The holes in NiO are consumed by methanol and photogenerated electrons accumulated on the conduction band of RP which participate in H_2 generation. This material also exhibited H_2 evolution activity in the near IR region.²⁰¹

Yan *et al.*²⁰² prepared nanocomposites consisting of CdS and amorphous MoO_x and studied photocatalytic hydrogen production under visible light irradiation. A 16 times enhancement in hydrogen evolution rates was observed compared to that of pristine CdS. The amounts of hydrogen evolved remained stable for five cycles. The photogenerated electrons from CdS resulted in an *in situ* reduction of Mo^{6+} in amorphous MoO_x creating electronic structure reconstruction in the nanocomposite.²⁰²



Chen *et al.*²⁰³ synthesized $g\text{-C}_3\text{N}_4$ modified with different amounts of Cu_2O nanoparticles by a simple calcination method. In the presence of Cu_2O , up to a 70% increase in visible light induced photocatalytic H_2 evolution was observed. Cu_2O nanoparticles acted as visible light sensitizers, induced sufficient charge separation and exhibited a negative shading effect so that their content above an optimum level reduced the number of active sites. The rate of hydrogen evolution remained stable for five cycles and the chemical state of Cu in the recovered catalyst was the same as that in the fresh one.²⁰³

Zhu *et al.*²⁰⁴ reported the use of wheat grain like textured TiO_2/CuO composite nanofibers synthesized by electrospinning for photocatalytic water splitting. CuO incorporation was found to enhance H_2 generation by about 16.8 times in comparison with that of pure TiO_2 . Significant loss of photocatalytic performance was not observed for four cycles.²⁰⁴

Moon *et al.*¹⁹¹ reported that size tunable Cu_2O nanodots decorated on the surface of TiO_2 nanocrystals got reduced to Cu

under UV irradiation. The resultant metal–semiconductor Schottky junction enhanced the hydrogen production rate significantly in comparison with P25 and reached approximately 50% of the one achieved with Pt/ TiO_2 counterparts, as shown in Fig. 16.¹⁹¹

Sadanandam *et al.*²⁰⁵ hydrothermally synthesized TiO_2 nanosheets with [001] facets exposed and impregnated with CuO quantum dots. It was observed that during the reaction CuO changed to Cu_2O and then to Cu. The energy band alignment of the double heterojunction $\text{TiO}_2/\text{Cu}_2\text{O}/\text{Cu}$ provided effective charge separation, increasing the H_2 production efficiency. Under visible light irradiation, the hydrogen production rate of the optimized material was approximately 10 times higher than that of pure TiO_2 , with an AQY of 26.93%. Cycling tests were carried out for the optimum catalyst and a H_2 production retention ratio of 96% is obtained after four cycles. The system was then purged with O_2 for 2 h to regenerate the used catalyst. After this, the retention ratio increased to 98%, indicating good cycling stability and photo corrosion resistance.²⁰⁵ The synergy between CuO and NiO, the most studied oxides as cocatalysts, was verified by Ravi *et al.*¹⁹² who synthesized $\text{CuO}@/\text{NiO}$ core@shell nanoparticles deposited on anatase TiO_2 and studied them for photocatalytic H_2 evolution under direct solar light irradiation. The rate of hydrogen evolution was found to be influenced by the thickness of the NiO shell and cocatalyst loading. 6-fold higher values were obtained in comparison with those of pure TiO_2 . The performance of this material was examined for few days. The setup was kept under dark conditions after 4 h solar light illumination and was purged with N_2 gas before the experiments on the next day. The rate of H_2 evolution remained similar for the first 2 days while a minor decrease was noticed on the third day. The authors attributed this to the decrease in the concentration of the sacrificial agent.¹⁹²

Another MO that was studied as a cocatalyst for water splitting is CoO. Mao *et al.*²⁰⁶ studied the photocatalytic behaviour of $g\text{-C}_3\text{N}_4/\text{CoO}$ nanocomposites with varying amounts of CoO, under visible light irradiation. This material with 0.5 wt% CoO content exhibited 3 times superior H_2 evolution rates compared to pure $g\text{-C}_3\text{N}_4$, owing to enhanced light absorption and separation of charge carriers. Stable performance was observed during three cycling experiments. Also, the recycled material exhibited no sign of aggregation.²⁰⁶

Zhu *et al.*²⁰⁷ prepared CoO_x confined in $g\text{-C}_3\text{N}_4$. The heterojunction formed primarily depended on the conditions of temperature treatment. The type II heterojunction formed between CoO nanoparticles and C_3N_4 nanotubes, under vacuum, exhibited the highest H_2 yield, due to the rapid electron transfer between CoO and C_3N_4 . $\text{CoO}/\text{C}_3\text{N}_4$ NTs were found to exhibit 6% activity loss while for $\text{Co}_3\text{O}_4/\text{C}_3\text{N}_4$ NTs it was almost 18.0%. A slight decrease in the intensity of the main peaks indexed to Co_3O_4 NPs was detected by XRD. This indicates the loss of a portion of anchored Co_3O_4 nanoparticles on C_3N_4 NTs during the recycling process. Also, some extent of agglomeration was observed for this material.²⁰⁷

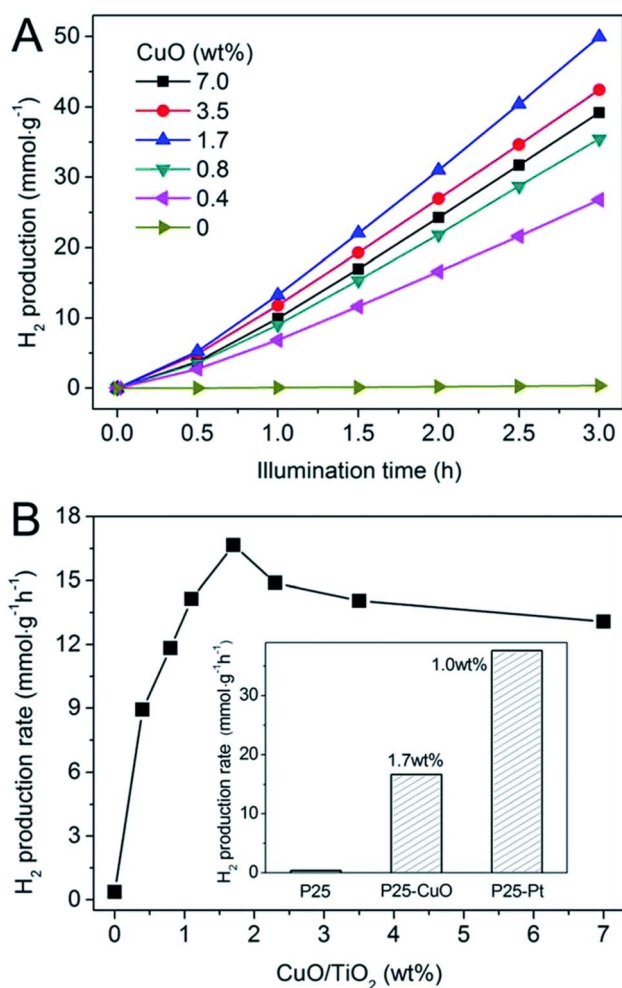


Fig. 16 (A) Hydrogen production vs. illumination time in the presence of CuO-decorated TiO_2 catalysts. (B) H_2 production rate vs. loading amount of CuO. The inset of (B) is a bar graph showing the activity of CuO-decorated TiO_2 in comparison to that of a Pt-decorated system. Reproduced with permission from ref. 191, Copyright 2014 The Royal Society of Chemistry.



4.3. Photocatalytic applications for CO₂ reduction

For photocatalytic conversion of CO₂, enhancing surface adsorption of CO₂ and suppressing charge carrier recombination represent one of the major focuses. As in the case of water splitting, transition MOs based on Cu, Ni, Co and Fe were most widely investigated as cocatalysts, along with CaO, MgO and In₂O₃ (see Table 5).

Uzunova *et al.*²⁰⁸ used DFT to deduce the mechanism of CO₂ photoconversion to methanol, using H₂O as the hydrogen source. The presence of MOs results in bending of a CO₂ molecule prior to hydrogenation and the bridging hydroxyl groups which resulted from water dissociation provided H₂. Methanol is formed by following a step-wise mechanism *via* the formation of carboxyl groups, formic acid and formaldehyde intermediates. A comparatively stronger interaction between CO₂ and Cu₂O compared to that between metallic catalysts was reported.²⁰⁸

Zhu *et al.*²⁰⁹ deposited Cu₂O clusters on TiO₂ nanosheets by photodeposition. The authors noticed the formation of surface oxygen vacancies on TiO₂ which was attributed to the presence of Cu₂O. Under simulated solar light irradiation and in the presence of water as the hydrogen source, methane was the only product formed. The presence of 1 wt% Cu₂O resulted in a 30 times enhancement of the photoactivity of pure TiO₂. Cu₂O clusters increased the surface alkalinity and enhanced CO₂ adsorption ability. High concentration of surface oxygen vacancies has also played the role of active sites for CO₂ conversion.²⁰⁹

Cu₂O loaded SiC photocatalysts were found to be selective towards methanol under visible light irradiation, using water as the hydrogen source. The CH₃OH yields were comparatively higher than that by the individual components.²¹⁰ In agreement with the report by Zhu *et al.*,²⁰⁹ Cu₂O nanoparticles electrochemically deposited on TiO₂ nanotubes selectively produced CH₄. Under visible light irradiation in the presence of water vapour, an optimum amount of octahedral Cu₂O nanoparticles exhibited high efficiency and selectivity towards methane.

The nanotube structure of TiO₂ provided active sites for the adsorption of reactants and facilitated the transport of photo-generated charge carriers. Synergistically, Cu₂O participated in light absorption and prevented oxidation of the reaction products.²¹¹

Fang *et al.*¹⁹⁴ synthesized CuO incorporated TiO₂ hollow microspheres with a hollow core and mesoporous shell structure, by a template free one pot HTT, followed by calcination. Photo-reduction of CO₂ was carried out over this material using H₂O as the hydrogen source under UV irradiation, producing CO and CH₄, as shown in Fig. 17. For an optimum CuO loading (3 wt%), 5.8- and 2.7-times higher CO and CH₄ yields, respectively, were obtained compared with those of pristine TiO₂ hollow microspheres, and 7.3- and 9.0-times higher yields were obtained compared to those of commercial TiO₂, respectively. The used catalysts were recovered from the reactor and stored under ambient conditions for 24 h. When this material was tested again, a minimum decrease in product yield was

observed indicating regeneration of the catalyst due to desorption of the gas products from its surface during storage.¹⁹⁴

Gui *et al.*¹⁹⁵ prepared MWCNT@TiO₂ core@shell nanocomposites, impregnated with a series of binary oxides (FeO_x, CuO_x, NiO, CoO_x and ZnO). The efficiency of these materials for CO₂ photoreduction was studied under visible light irradiation in the presence of water vapour. Methane was selectively produced and CuO modified MWCNT@TiO₂ exhibited superior performance among the series. The better performance was attributed to the MO band position with respect to that of TiO₂.¹⁹⁵

Tahir *et al.*²¹² used cordierite honeycomb monoliths loaded with N/TiO₂ and Cu/TiO₂ nanocatalysts using a sol-gel dip coating method for dynamic photocatalytic CO₂ reduction with H₂ selectively to CO under UV-light irradiation. Both Cu and N doping resulted in a shift of band gap energy towards the visible region and also in an efficient transport and trapping of electrons, hindering carrier recombination. The presence of Cu resulted in a 14 times higher CO production than N/TiO₂ and 64 times higher CO production than that of pure TiO₂. Here, the CO production decreased after 4 h of irradiation time, indicating deterioration of photoactivity.²¹²

Shoji *et al.*²¹³ loaded Cu_xO as a reduction cocatalyst and cobalt phosphate (CoPi) as an oxidation cocatalyst on SrTiO₃ nanorod thin films. Photocatalytic experiments under UV irradiation were conducted by immersing the film in an aqueous solution bubbled with CO₂. Selective evolution of CO was observed with the highest rate exhibited by the materials with dual cocatalysts. The CO generation rate was found to increase linearly in accordance with the irradiation time. Here, Cu_xO nanoclusters facilitated CO₂ reduction whereas CoPi facilitated surface oxidation to promote oxygen generation. The dual cocatalyst incorporated materials exhibited a 2.5 times superior CO generation rate compared to pure SrTiO₃.²¹³

Junior *et al.*²¹⁴ synthesized layered semiconductor K₂Ti₄O₉ by the solid-state method and pillared it with TiO₂. The resultant mesoporous composite was then loaded with Cu₂O nanoparticles as the cocatalyst. CO₂ photoreduction was carried out using water as the hydrogen source under simulated solar light. Improved performance was observed for the pillared material TiO₂/Ti₄O₉ than the pristine layered potassium tetratitanate. Here, electron transfer between the guest TiO₂ and the host Ti₄O₉²⁻ layers could have played a role, along with the presence of additional photoactive sites. Loading of Cu₂O nanoparticles resulted in further improvement of photo activity due to enhanced visible light harvest promoting electron transfer with TiO₂ and Ti₄O₉²⁻. It is to be noted here that since the temperature in the system reached 70 °C during the reaction, the contribution of photothermal effects cannot be ignored.²¹⁴

Jeyalakshmi *et al.*²¹⁵ hydrothermally synthesized Na_(1-x)La_xTaO_(3+x) and Pt, Ag, Au, CuO, NiO and RuO₂ were introduced as cocatalysts by following the wet impregnation method. CO₂ photoconversion was carried out in an alkaline medium under UV-visible light. Methanol and ethanol were produced as the major products along with traces of methane, ethane, ethylene, *etc.* The amounts of methanol produced remained consistent for three cycles over NiO/Na_(1-x)La_xTaO_(3+x). High AQYs among



Table 5 Compilation of synthetic and photocatalytic aspects of MO-based materials in CO₂ conversion^a

Cocatalyst/ photocatalyst	Synthetic method	Reactants	Light region	Products [$\mu\text{mol g}^{-1} \text{h}^{-1}$]	AQY [%]	Ref.
Cu ₂ O/TiO ₂ nanosheets	Photodeposition and calcination in air (200 °C, 2 h)	CO ₂ and H ₂ O	Simulated solar light	CH ₄ (8.68)	NP	209
Cu ₂ O/SiC	Precipitation using NaOH followed by chemical reduction using N ₂ H ₄ ·H ₂ O	NaOH, Na ₂ SO ₃ and CO ₂	Visible	CH ₃ OH (38)	NP	210
Octahedral Cu ₂ O/TiO ₂ nanotube arrays	Electrochemical deposition	CO ₂ and H ₂ O	Visible	CH ₄ (~400 ppm for 4 h)	NP	211
CuO–TiO ₂ hollow microspheres	HTT (180 °C for 20 h)	CO ₂ and H ₂ O	UV	CO (14.5) CH ₄ (2.1) H ₂ (2.8)	1.285 0.747	194
FeOx-MWCNT@TiO ₂	Impregnation and calcination under N ₂ (400 °C for 2 h)	CO ₂ and H ₂ O	Visible	CH ₄ (0.85 $\mu\text{mol g}^{-1}$ for 8 h)	NP	195
CuO-MWCNT@TiO ₂	Impregnation and calcination under N ₂ (400 °C for 2 h)	CO ₂ and H ₂ O	Visible	CH ₄ (0.93 $\mu\text{mol g}^{-1}$ for 8 h)	NP	195
NiO-MWCNT@TiO ₂	Impregnation and calcination under N ₂ (400 °C for 2 h)	CO ₂ and H ₂ O	Visible	CH ₄ (0.44 $\mu\text{mol g}^{-1}$ for 8 h)	NP	195
CoO-MWCNT@TiO ₂	Impregnation and calcination under N ₂ (400 °C for 2 h)	CO ₂ and H ₂ O	Visible	CH ₄ (0.38 $\mu\text{mol g}^{-1}$ for 8 h)	NP	195
ZnO-MWCNT@TiO ₂	Impregnation and calcination under N ₂ (400 °C for 2 h)	CO ₂ and H ₂ O	Visible	CH ₄ (0.27 $\mu\text{mol g}^{-1}$ for 8 h)	NP	195
Cu–TiO ₂ in a cordierite honeycomb monolith	Dip coating and calcination in air (500 °C for 5 h)	CO ₂ and H ₂	Simulated solar light	CO (763)	0.0826	212
Cu _x O/CoPi–SrTiO ₃ nanorod thin films	Impregnation or photoelectrochemical deposition	CO ₂ and H ₂ O	UV	CH ₄ (4.20)	NP	213
Cu ₂ O on TiO ₂ -pillared K ₂ Ti ₄ O ₉	Calcination in air (800 °C for 20 h), ion-exchange (using HCl), exfoliation using (TBAOH), impregnation and reflux (160 °C for 2 h)	CO ₂ and H ₂ O	Simulated solar light	CH ₃ OH (2.93 $\mu\text{mol g}^{-1}$ for 5 h)	NP	214
CuO/Na _(1-x) LaxTaO _(3+x)	Impregnation and calcination in air (300 °C for 2 h)	NaOH, CO ₂ and H ₂ O	UV-visible	CH ₃ OH (60.5) C ₂ H ₅ OH (15.9) C ₂ H ₄ O (0.9) C ₃ H ₆ (0.15) C ₂ H ₆ (0.04) C ₂ H ₄ (0.03) CH ₄ (0.01) H ₂ (0.2)	0.0105	215
NiO/Na _(1-x) LaxTaO _(3+x)	Impregnation and calcination in air (270 °C for 2 h)	NaOH, CO ₂ and H ₂ O	UV-visible	CH ₃ OH (59.6) C ₂ H ₅ OH (18.8) C ₂ H ₄ O (2.15) C ₃ H ₆ (0.4) C ₂ H ₆ (0.01) C ₂ H ₄ (0.03) CH ₄ (0.02) H ₂ (0.2)	0.0112	215
Co ₃ O ₄ /K ₂ Ti ₆ O ₁₃	Calcination in air (800 °C for 12 h)	CO ₂ and H ₂ O	Visible	CH ₃ OH (36.4 $\mu\text{mol g}^{-1}$ for 3 h) HCHO (453.2 $\mu\text{mol g}^{-1}$ for 3 h) CH ₄ (1.3 $\mu\text{mol g}^{-1}$ for 3 h) H ₂ (8 $\mu\text{mol g}^{-1}$ for 3 h)	NP	216
NiO/K ₂ Ti ₆ O ₁₃	Calcination in air (800 °C for 12 h)	CO ₂ and H ₂ O	Visible	CH ₃ OH (31.2 $\mu\text{mol g}^{-1}$ for 3 h) HCHO (99.7 $\mu\text{mol g}^{-1}$ for 3 h) CH ₄ (1.1 $\mu\text{mol g}^{-1}$ for 3 h) H ₂ (201 $\mu\text{mol g}^{-1}$ for 3 h)	NP	216



Table 5 (Contd.)

Cocatalyst/ photocatalyst	Synthetic method	Reactants	Light region	Products [$\mu\text{mol g}^{-1} \text{h}^{-1}$]	AQY [%]	Ref.
CuO/ $\text{K}_2\text{Ti}_6\text{O}_{13}$	Calcination in air (800 °C for 12 h)	CO_2 and H_2O	Visible	CH_3OH (159.7 $\mu\text{mol g}^{-1}$ for 3 h) HCHO (41.4 $\mu\text{mol g}^{-1}$ for 3 h) CH_4 (0.8 $\mu\text{mol g}^{-1}$ for 3 h) H_2 (282 $\mu\text{mol g}^{-1}$ for 3 h)	NP	216
Cu- $\text{Bi}_2\text{O}_3/\text{TiO}_2$	Calcination under H_2 (550 °C for 5 h)	CO_2 and H_2O	UV-visible	CO (5.74) CH_4 (11.90)	NP	217
NiO/ InTaO_4	Sol-gel method and calcination in air (1100 °C for 12 h)	NaOH , CO_2 and H_2O	Sunlight	CH_3OH (11.3)	0.063	218
NiO- InNbO_4	Incipient wetness impregnation and calcination in air (350 °C for 1 h)	CO_2 and H_2O	Simulated solar light	CH_3OH (1.577)	NP	219
Co_3O_4 - InNbO_4	Incipient wetness impregnation and calcination in air (350 °C for 1 h)	CO_2 and H_2O	Simulated solar light	CH_3OH (1.503)	NP	219
NiO- KTaO_3	HTT (180 °C for 12 h) and calcination in air (600 °C for 1 h)	CO_2 and isopropanol	UV	CH_3OH (1523)	NP	220
NiO- InVO_4	Impregnation and calcination in air (350 °C for 2 h)	CO_2 , H_2O and KHCO_3	Visible	CH_3OH (1.526)	NP	221
Ni@NiO/ InTaO_4 -N	Impregnation, chemical reduction (using NaBH_4) and calcination in air (200 °C for 1 h)	CO_2 and H_2O	Visible	CH_3OH ($\sim 300 \mu\text{mol g}^{-1}$)	NP	222
Macro/microporous MgO modified TiO_2	Biotemplating and calcination in air (500 °C for 3 h)	CO_2 and H_2O	Simulated solar light	CH_4 (1.87)	NP	223
TiO_2/MgO	HTT (200 °C for 12 h)	CO_2 , H_2O and NaOH	UV	CO (70.7 $\mu\text{mol g}^{-1} \text{L}^{-1}$ for 24 h) CH_4 (34.5 $\mu\text{mol g}^{-1} \text{L}^{-1}$ for 24 h) HCOOH (19.5 $\mu\text{mol g}^{-1} \text{L}^{-1}$ for 24 h) CH_3COOH (24.9 $\mu\text{mol g}^{-1} \text{L}^{-1}$ for 24 h) CH_4 (82 $\mu\text{mol g}^{-1} \text{L}^{-1}$ for 8 h)	NP	224
Ca/ $\text{Ti}_x\text{SiMCM-41}$	HTT (100 °C for 12 h) and calcination (500 °C for 2 h)	CO_2 and H_2O	UV	CH_4 (82 $\mu\text{mol g}^{-1} \text{L}^{-1}$ for 8 h)	NP	225

^a ~ Data taken from a graphical representation of the corresponding reference; NP: value not provided.

the series were obtained for 0.2 wt% NiO (11.29%) and 1 wt% CuO (10.50%) as cocatalysts. These oxides reduced the band gap energy of NaTaO_3 and at the same time the light absorption onset edge was found to be extended to the visible region.²¹⁵

Garay-Rodriguez *et al.*²¹⁶ used an ultrasound assisted sol-gel method for synthesizing MO (Co_3O_4 , NiO, and CuO) incorporated $\text{K}_2\text{Ti}_6\text{O}_{13}$. Interestingly while some extent of Co^{2+} cations were found to selectively substitute Ti^{4+} cations, Ni^{2+} and Cu^{2+} were incorporated in both Ti^{4+} and K^+ sites. The metal-cation introduction between the tunnels favoured the growth of potassium-poor phases as impurities, especially in the CuO loaded $\text{K}_2\text{Ti}_6\text{O}_{13}$ samples. Photoconversion of CO_2 using H_2O as the hydrogen source under UV-visible light, resulted in different products depending on the loaded MO. It was observed that while the presence of Co_3O_4 favoured the formation of formaldehyde, NiO favoured hydrogen evolution due to additional photo-reforming of the organic compounds and CuO enhanced the selectivity towards methanol and hydrogen. The product

yields remained consistent for three consecutive reaction cycles for Co_3O_4 , NiO, and CuO incorporated $\text{K}_2\text{Ti}_6\text{O}_{13}$. Moreover, no structural changes were observed by comparing the XRD patterns of the fresh and used catalysts.²¹⁶

Jeong *et al.*²¹⁷ explored the use of Bi_2O_3 as a promoter for Cu cocatalysts anchored on the surface of TiO_2 nanoparticles for the photocatalytic conversion of CO_2 to CH_4 , under UV-visible light irradiation. The presence of Bi_2O_3 increased the product selectivity towards methane (see Fig. 18). In the absence of Bi_2O_3 , the photogenerated electrons reduce the CO_2 adsorbed on the cocatalyst surface to CO, which has a low binding strength and will get detached as the product. Bi_2O_3 provided additional binding sites for CO where it undergoes further transformation, producing CH_4 .²¹⁷

Wang *et al.*²¹⁸ used a NiO loaded InTaO_4 photocatalyst to carry out both aqueous phase and vapor phase photoconversion of CO_2 using visible light. The rate of methanol production was $11.1 \mu\text{mol g}^{-1} \text{h}^{-1}$ which was slightly increased up to $11.30 \mu\text{mol}$



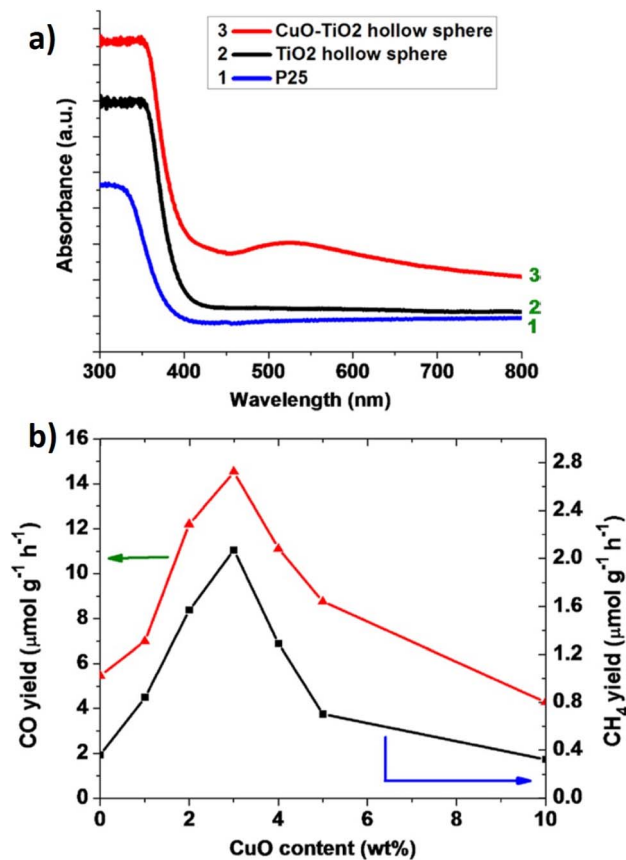


Fig. 17 (a) UV-vis DRS for various TiO₂ based nano-materials. The CuO content in the CuO-TiO₂ catalyst is 3 wt%. (b) Yields of the gas products (CO and CH₄) produced during 24 h of photo-driven CO₂ reduction on the TiO₂ hollow microsphere catalysts with various Cu contents. Reproduced with permission from ref. 194, Copyright 2015 American Chemical Society.

g⁻¹ h⁻¹ by utilizing concentrated sunlight. The quantum efficiencies were estimated to be 0.0045% and 0.063% in aqueous-phase and gas phase reactors, respectively.²¹⁸

Lee *et al.*²¹⁹ synthesized InNbO₄ from the corresponding binary oxides, and cocatalysts (NiO and Co₃O₄) were impregnated by following the incipient-wetness method. NiO-InNbO₄ after a reduction-oxidation pre-treatment exhibited the highest activity for producing methanol under visible light which was attributed to the presence of core-shell type Ni⁰/NiO on the surface and the presence of a small amount of Nb₂O₅ as a promoter.²¹⁹

Comparative studies were carried out between hydrothermally synthesized perovskite-type and pyrochlore-type powders of potassium tantalate (KTaO₃). The pyrochlore form exhibited better photocatalytic activities for producing methanol under UV light irradiation which was further increased by adding NiO as a co-catalyst.²²⁰ Lee *et al.*²²¹ synthesized InVO₄ by the solid-state reaction method using In₂O₃ and V₂O₅ as the starting materials and NiO was added as the cocatalyst by incipient wetness impregnation. The photocatalytic reduction of CO₂ with water was carried out under visible light illumination, producing methanol. Introduction of NiO resulted in lowering

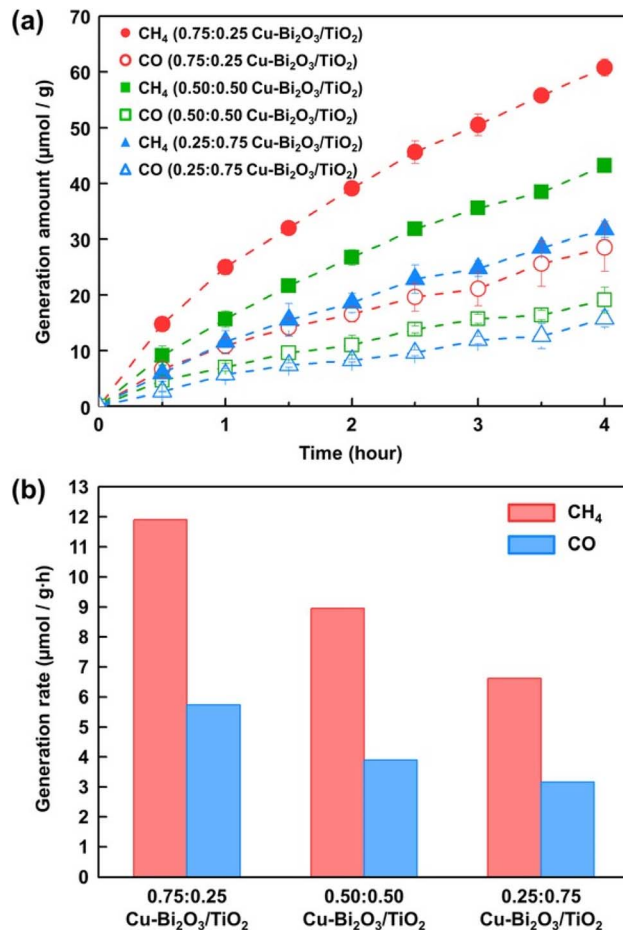


Fig. 18 (a) Amount of CH₄ and CO generated upon using 0.75 : 0.25, 0.50 : 0.50, and 0.25 : 0.75 Cu-Bi₂O₃/TiO₂ photocatalysts. (b) Rates of CH₄ and CO generation in the first 4 h of irradiation. Reproduced with permission from ref. 217, Copyright 2016 European Chemical Societies Publishing.

of the bandgap due to the formation of sub bands in the valence band by creating additional defect sites on the surface of InVO₄, resulting in a higher methanol yield with an AQY of 2.8%.²²¹

Tsai *et al.*²²² synthesized N-doped InTaO₄ photocatalysts and introduced Ni@NiO core@shell nanostructures as a cocatalyst for transforming CO₂ into methanol. Under visible light irradiation, a 2-fold increment in methanol yield was observed for the N-doped samples which was further increased by the presence of Ni@NiO cocatalysts, compared to that of pristine InTaO₄. Due to a narrow band gap, Ni@NiO/InTaO₄-N absorbed more light, and the Ni@NiO core@shell structures promoted more efficient electron transport.²²²

Wang *et al.*²²³ synthesized porous MgO modified TiO₂ using water convolvulus as the biotemplate. This material photoreduces CO₂ in the presence of water vapour, selectively producing methane. 0.2 wt% MgO resulted in a 3.5 times increase in the CH₄ evolution rate in comparison with pure TiO₂. The authors attributed this to various factors including CO₂ adsorption capacity, charge carrier transport, and enhanced number of active sites, which depend on the amount of added MgO.²²³



Table 6 Compilation of synthetic and photocatalytic aspects of MO-based materials in the NRR

Cocatalyst/p hotocatalyst	Preparation methods	Reaction conditions	Light source	Sacrificial reagent	NH ₃ production ^a	AQY [%]	Ref.
Bi ₂ MoO ₆ /OV-BiOBr	HTT (120 °C for 8 h)	30 mg of the photocatalyst in 60 mL of ultrapure water, RT	300 W xenon lamp	None	90.7 μmol g ⁻¹ h ⁻¹	NP	226
30 wt% SV-1T-MoS ₂ /CdS	HTT (200 °C for 24 h)	20 mg of photocatalyst in 100 mL 20 vol% methanol solution	Simulated solar light	Methanol	8220.83 μmol L ⁻¹ g ⁻¹ h ⁻¹	4.424	227
18% Sb ₂ O ₃ /W ₁₈ O ₄₉	Solvothermal (180 °C for 20 h)	5 mg photocatalyst in the aqueous solution (10 mL) containing 20 vol% methanol, RT	300 W xenon lamp	Methanol	35.3 μmol g ⁻¹ h ⁻¹	0.063	228
g-C ₃ N ₄ /40 wt% Fe ₂ O ₃	Ternary deep eutectic solvents and calcination (550 °C for 4 h)	10 mg of photocatalyst in 100 mL of distilled water, RT	300 W xenon lamp	Methanol	4380 μmol L ⁻¹ h ⁻¹	NP	229

^a The production rates were expressed as how they appear in the references.

Torres *et al.*²²⁴ mixed MgO with TiO₂ in different proportions and the resultant nanocomposites were studied in the photoconversion of CO₂. Enhanced selectivity was observed for CO, while CH₄, HCOOH and CH₃COOH were also produced in lower amounts. Addition of 1 wt% of MgO enhanced CH₄ production compared to that with pure TiO₂, however, a further increase in the MgO content significantly deteriorated the performance. The authors attributed this effect to the insulating behaviour of MgO. It is also likely that higher amounts of MgO are blocking the active sites, thereby lowering the light absorption properties.²²⁴

Jo *et al.*²²⁵ used a series of MCM-41 mesoporous silica modified with Ca and Ti for the photoreduction of CO₂ to CH₄. The optimum material among the series exhibited extremely high yields of methane under solar light irradiation. Optimum Ca doping contributed towards photoreduction by providing additional sites for CO₂ adsorption and by promoting charge carrier separation. It is to be noted that excess metal sites diminish the performance by providing charge recombination centers.²²⁵

4.4. Photocatalytic applications for N₂ photofixation

In addition to the well-known photocatalytic applications previously mentioned, MOs are also utilized as a cocatalyst in photocatalytic systems for the photoreduction of N₂ to produce NH₃. Cocatalysts as MOs are used to broaden the range of absorbed light, as well as to facilitate the electron transport throughout the system and stop electron-hole recombination. Table 6 summarizes several examples of MO-based photocatalysts used in recent years for the NRR.

For enhanced photocatalytic performance for N₂ conversion to ammonia without any noble metal and sacrificial agents under ambient conditions, Xue *et al.*²²⁶ produced heterojunctions based on n-type Bi₂MoO₆ nanorods and oxygen-vacancy-rich p-type BiOBr nanosheets (Bi₂MoO₆/OV-BiOBr). The average ammonia generation rate of the Bi₂MoO₆/OV-

BiOBr heterojunctions is 90.7 μmol g⁻¹ h⁻¹, which is about 30 times higher than that of pristine Bi₂MoO₆ nanorods (3.0 μmol g⁻¹ h⁻¹) and 3 times higher than that of OV-BiOBr nanosheets (31.2 μmol g⁻¹ h⁻¹). This study highlights the significantly increased photocatalytic N₂ fixation activity of the Bi₂MoO₆/OV-BiOBr heterojunctions, without the use of any sacrificial agents.

Sun *et al.*²²⁷ reported sulfur vacancy (SV)-rich O-doped 1T-MoS₂ nanosheets (denoted as SV-1T-MoS₂), prepared by HTT, as cocatalysts to assist in the photocatalytic nitrogen fixation of CdS nanorods, as a hybrid material SV-1T-MoS₂/CdS. Different quantities of the cocatalyst, ranging from 5 to 100 mg, representing 5 to 100 wt%, were used to produce the SV-1T-MoS₂/CdS composites. A remarkable photocatalytic N₂ reduction rate of 8220 μmol L⁻¹ h⁻¹ g⁻¹ is demonstrated by the optimized 30 wt% SV-1T-MoS₂/CdS composites, under simulated solar light irradiation (AM 1.5 G), with methanol as a scavenger, obtaining an AQE of 4.424%. This rate is 2.4 and 4.6 times greater than that of CdSPT (0.1 wt%) and CdS nanorods, respectively, suggesting that SV-1T-MoS₂ may exceed Pt in the CdS's photocatalytic nitrogen fixation performance. However, the rate of NH₄⁺ will decrease as the amount of SV-1T-MoS₂ in the composites increased from 30 to 100 wt%. This is probably because excessive MoS₂ prevents the light from being used. Meanwhile, under visible light (780 nm > λ > 420 nm), the photocatalytic nitrogen fixation rate was 3168 μmol L⁻¹ h⁻¹ g⁻¹.

Three cycles were performed to evaluate the stability over 30 wt% SV-1T-MoS₂/CdS, and after 9 hours of testing, 80% of the incipient activity was still present, demonstrating the materials' good cycle stability.

Sb₂O₃/W₁₈O₄₉, a composite made of 2D Sb₂O₃ nanosheets and 1D W₁₈O₄₉ nanowires, was created by Hui *et al.*²²⁸ via the solvothermal process, by varying the Sb₂O₃ weight, ranging from 1.8 to 70%. The highest NH₃ production was obtained for 18% Sb₂O₃/W₁₈O₄₉ with a production rate of 35.3 μmol g⁻¹ h⁻¹, with methanol as a scavenger under visible light illumination, which is 3.4 and 5.5 times higher than that of W₁₈O₄₉ and



Sb_2O_3 , respectively, with an AQE of 0.063%. However, as the Sb_2O_3 concentration was increased above 18 wt%, the rate of NH_3 synthesis decreased. This is probably due to Sb_2O_3 aggregation and the low $\text{Sb}_2\text{O}_3/\text{W}_{18}\text{O}_{49}$ interface active regions, which inhibited N_2 adsorption and activation. The heterostructures demonstrated good repeatability with approximately consistent NH_3 production rates. Additionally, cycling studies showed that even after five consecutive cycles, the catalytic activity was mainly the same, with a certain decrease of activity due to aggregation.

Mou *et al.*²²⁹ created $\text{g-C}_3\text{N}_4/\text{MO}$ composites utilizing a simple one-step calcination procedure and a ternary deep eutectic solvent (DES) technique, in which $\text{g-C}_3\text{N}_4/\text{Fe}_2\text{O}_3$ was of interest due to its photocatalytic response. Melamine was added in various amounts to control the loading of $\alpha\text{-Fe}_2\text{O}_3$, ranging from 0.1582 g to 2.291 g for the production of 50%-DES, 40%-DES, 35%-DES, and 30%-DES, respectively. At 40 wt%, the best activity was attained ($\text{g-C}_3\text{N}_4/40$ wt% Fe_2O_3) with an average rate of $4380 \mu\text{mol L}^{-1} \text{h}^{-1}$ for producing NH_3 , with methanol acting as a scavenger. The effective interfacial contact between 2D $\text{g-C}_3\text{N}_4$ nanosheets and $\alpha\text{-Fe}_2\text{O}_3$ nanoparticles, which improves the transfer and separation efficiency of the charge carriers, may be the reason for the high photocatalytic activity.

5. Comparison between MXenes and metal oxides

MXenes and MOs have gained a lot of interest and have been investigated for photocatalytic applications due to their unique properties, such as electronic band structures, catalytic edge effects, planar morphology, *etc.*²³⁰ This section is dedicated to a comparison as succinct as possible between these two categories of materials when they are used as cocatalysts in photocatalytic reactions. The most important criteria taken into account was the employed synthesis methods, their role in promoting photogenerated charge separation, environmental issues, the cost/benefit ratio, the number of combinations in which they can be used and their photocatalytic stability.

Thus, in terms of synthesis conditions, both categories of materials can be obtained by following relatively simple approaches. While chemical liquid-phase etching of the MAX phase results in the corresponding MXenes, MOs in their amorphous, polycrystalline or single crystal form can be synthesized from their precursors (nitrates, chlorides, *etc.*) under thermal or hydrothermal conditions.²³¹ In the previous sections, a summary of individual synthesis methods of the two categories of materials is provided. While making a comparison, we need to take into account the fact that MXenes are still in their infancy, whereas MOs have a long history of evolution. A vast majority of photocatalytic studies using MXenes were carried out using $\text{Ti}_3\text{C}_2\text{T}_x$ while for oxides several material options were available. For a fair and realistic assessment, the strategies frequently used to integrate these cocatalysts (MXenes and MOs) with the photocatalysts are to be considered. Thus, the choice of a method relies not only on maintaining the structural, compositional and morphological integrity of the

materials employed, but also on establishing a homogeneous and intimate contact between them. In addition, the amount of cocatalyst used is important since an excess can be detrimental for the overall performance.

Broadly, the methods used to develop cocatalyst-photocatalyst composites can be classified into two categories: (i) multi-step synthesis – the components can be synthesized separately and can be combined in desired proportions in a post-synthetic manner; and (ii) “one-pot” approaches – in which the cocatalyst and the photocatalyst are formed simultaneously under the same conditions. As mentioned in the previous sections, post-synthetic approaches are always required for MXenes, since the etching conditions (usually HF) are not compatible with the semiconductor photocatalysts. Also, in cases where the desired phase formation and thermal stabilization of the composites under oxidizing conditions are required, total destruction of MXenes can occur, forming the corresponding MOs. It is to be noted here that a few studies reported controlled oxidation of MXenes (as a precursor) producing cocatalyst-photocatalyst composites. However, it is difficult to control the cocatalyst-photocatalyst ratio in this way and to date, it is not sure whether this approach can be generalized. In our understanding, HTT is the most optimum method that can be adapted to both MXenes and MOs to produce photocatalyst composites. An advantage here for MXenes is their rich surface chemistry that enables them to bind appropriately with the semiconductors. Physical mixing approaches such as grinding are also compatible with both these families of materials; however, it is difficult to establish an intimate contact between the cocatalyst and photocatalyst as required to maintain their synergistic functioning.

In photocatalytic reactions, the main role of cocatalysts is to promote the separation of photogenerated charge carriers by acting either as electron mediators or hole mediators. In this regard, starting from their discovery, due to their excellent electrical conductivity (possessing mostly metallic or semi-metallic properties and thus mainly forming Schottky junctions in contact with the semiconductor photocatalysts),²³² MXenes have been mainly promoted as a promising reservoir to trap and shuttle the photogenerated electrons from semiconductors. However, as was demonstrated in many studies, the MXene's role of electron- or hole-mediator depends very much on the type of surface functional groups^{37,102,233–236} and/or the work function compatibility between the MXene and the semiconductor forming the composites.²³⁷ This latter situation is also applicable in the case of photocatalytic composites with MOs as cocatalysts. Consequently, across the interfaces between the components of the composite photocatalysts, photogenerated holes will flow from the high-work-function component (the semiconductor acting as a photocatalyst) to the low-work-function component (the MO acting as a cocatalyst), while the photogenerated electrons will follow the opposite direction. In other words, the MO cocatalyst, which is actually a semiconductor material, might work either as an electron mediator or a hole mediator depending on the type of junction it makes with the semiconductor photocatalyst (*e.g.*, p-type/n-type, n-type/n-type junction or even *Z* or *S* scheme). To

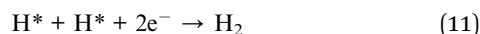
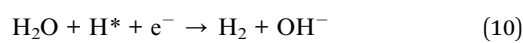


summarize, in terms of the type of junction formed within the composite photocatalyst, the main difference between the MXenes and the MOs is given by their intrinsic properties. MXenes generally possessing a metallic character form Schottky junctions with the semiconductor photocatalyst depending on the terminal functional groups (*e.g.*, $-\text{OH}$, $-\text{O}$, and $-\text{F}$). On the other hand, MOs generally form either p-type/n-type or n-type/n-type junctions as they have semiconductor properties.

Regarding the environmental concerns associated with the use of MXenes or MOs as cocatalysts that might arise, it should be noted that MXene synthesis is much more harmful to the environment than that of the MOs, mainly due to the use of the HF etchant. However, the use of proper handling and disposal measures can minimize the potential hazards, as well as the use of alternative synthetic methods that are currently being developed.

We will reach the same conclusion if we compare the production costs (the production costs for MXenes are comparatively higher, considering the synthesis of parent MAX phases), but perhaps the fairest cost-benefit comparison would be to follow the gain resulting from their use in the three types of reactions discussed in this review.

Regarding the photocatalytic water splitting reaction, multiple reactions occur on the surface of the composite photocatalyst as represented in eqn (9)–(11):



As shown, in the first step (eqn (9)), an intermediate adsorbed H^* is formed and then, in the second step, the H_2 is desorbed by two possible ways: the reaction between the adsorbed H^* and a H^+ (eqn (10)) or through the combination of two H^* (eqn (11)). In the evaluation of photoactivity of a composite, the Gibbs free energy of the adsorbed hydrogen (ΔG_{H^*}) is regarded as a crucial parameter, and in an ideal situation its value should be close to zero. A large negative or positive ΔG_{H^*} indicates too strong or too weak chemical adsorption of H^* , meaning a harder or easier desorption of the H_2 molecule as the reaction product, respectively. In this view, it has to be specified that MXenes show in general $|\Delta G_{\text{H}^*}|$ values closer to zero than the MOs. Thus, for example, the O-terminated Ti_3C_2 shows a near-zero $|\Delta G_{\text{H}^*}|$ value (0.00283 eV),¹¹¹ which is even lower than the values of noble metal platinum (0.09 eV),^{238,239} MoS_2 (0.08 eV)²⁴⁰ and WS_2 (0.22 eV),²⁴⁰ which are among the most active cocatalysts for the water splitting reaction, while TiO_2 shows ΔG_{H^*} values of 2.37 eV and 2.32 eV for the (101) and (001) exposed facets,²⁴¹ respectively. Actually, this could be an explanation for the very good photoactivity observed for the MXene-based composites.

In the case of both photocatalytic CO_2 reduction and N_2 photofixation reactions, the common step is the activation of the reactants (*i.e.*, CO_2 and N_2). As both reactants are acid molecules (with CO_2 possessing much higher acidity than N_2), the presence of basic sites, such as the hydroxyl groups, is

a prerequisite to perform the activation of these molecules. In view of that, in the case of MXenes, surface alkanisation is needed in order to create more surface hydroxyl groups as active sites¹³¹ to adsorb and activate both CO_2 and N_2 molecules. This post-synthetic procedure is not necessary in the case of MOs, which already possess large amounts of such hydroxyl groups on the surface.

To summarize, in terms of the best activities reached so far, studies on water splitting indicate that a combination of MXenes ($\text{Ti}_3\text{C}_2\text{T}_x$) with black phosphorous and g- C_3N_4 exhibited the best performance for hydrogen evolution. Also, superior performances were observed for MXenes combined with CdS. In addition, TEA and lactic acid were reported to be the most appropriate hole scavengers under visible light irradiation. In the case of MOs, the highest H_2 production rates were observed for CuO as the cocatalyst in the presence of methanol or glycerol as hole scavengers. It is clear from the literature data that extremely high values for AQY were obtained when MXenes were used as cocatalysts. On the other hand, for photocatalytic CO_2 conversion, selectivity is a major parameter to consider. Using MXenes as the cocatalyst, CO and CH_4 were found to be the main products in most of the cases. Even though a few studies reported the formation of methanol, the product window was narrow. Instead, as a general trend, MOs were found to be more selective for hydrocarbons (other than CH_4) and oxygenated products like methanol. There are no big differences in the photocatalytic activity when MOs or MXenes are used as the cocatalyst for N_2 photofixation, because the product formation and selectivity are related to the cocatalyst-photocatalyst composite, rather than the individual components.

Regarding the number of possible combinations, by comparing Tables 1–3 with Tables 4–6 one can observe that in the case of MXenes there are limited combinations used, mainly Ti-based MXenes, while for the MOs many more possibilities were explored.

As the last comparison criterion, the durability of the composites based on MXenes and metal oxides represents an important aspect that has to be considered for practical applications. Thus, it has to be specified that leaving a photocatalyst for a longer period of time under the harsh conditions imposed by the photocatalytic reactions will finally lead to photocorrosion phenomena.²⁴² In this view, in comparison with the MOs, one of the biggest advantages of using MXenes as cocatalysts is to improve the photostability of the composites that contain it by shuttling photogenerated carriers from the unwanted side reaction with semiconductors.^{243,244}

To conclude, regardless of the comparison criteria used here, both categories of materials present advantages that should further guide research in the field of photocatalysis towards finding the most efficient cocatalyst/photocatalyst combinations (see Fig. 19).

6. Summary and future perspectives

This review underlined how important is to optimize the amount and type of the cocatalyst, and its choice should be



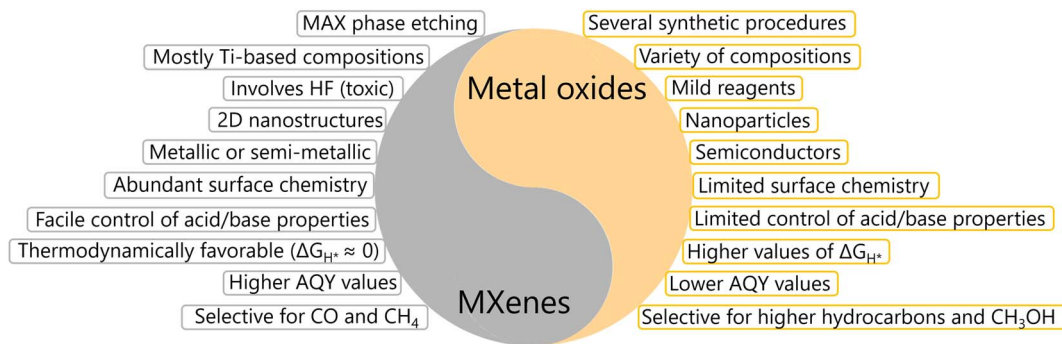


Fig. 19 Scheme summarising the advantages and disadvantages of MXenes and metal oxides to be used as cocatalysts in photocatalytic reactions.

carefully addressed and controlled. Moreover, the electronic properties, composition, loading, and morphology, among other factors, play an important role, affecting the overall photocatalytic activity of the hybrid materials. The literature survey presented in this review demonstrated that the use of MXenes and MOs as cocatalysts could greatly improve the photoactivity of a photocatalyst. This is mainly due to their capacity to promote the separation of photogenerated charge carriers. Overall, the photocatalytic processes presented in this review appear to be very attractive; however, there are challenges, such as activity and stability or high cost, that need to be elucidated.

Since the photocatalytic process takes place on the surface of the photocatalysts, the screening of the surface and electronic properties of the photocatalytic composites should be more systematic. This is mandatory especially in the case of MXenes as their surface is terminated with different functional groups, the type of which not only depending on the etching procedure but also altering under reaction conditions, thus changing the photocatalytic behaviour of the MXene-based composites. In this view, a single characterization approach (which in most cases is carried out before performing the photocatalytic reaction) is insufficient to support a comprehensive understanding of the surface properties of MXenes and thus, a combination of complementary characterization techniques is required.

With regard to the stability issues of the currently used photocatalytic systems, it should be underlined that the effects of the reaction environment on the composite's surface demand more research attention. On this subject, as compared with MOs, MXenes have not been as extensively investigated as cocatalysts in photocatalytic processes. As already mentioned, MXenes are unstable in aqueous environments and in the presence of O_2 and UV irradiation, because under such conditions MXenes are transformed into their corresponding oxides.⁴⁵ However, as shown within this review in subSections 3.3.1 and 3.4.1, this can result in a positive approach for obtaining materials with properties of both type of materials, carbides and oxides, paving the way for new materials for photocatalysis. Moreover, there are studies that confirm the role of carbon in facilitating the separation of charge carriers and the use of MXenes as a precursor for photocatalysts.^{105,158,159} To sum up, in regard to the stability of MXenes and MOs acting as

cocatalysts, probing approaches at the molecular level, such as different *in situ/operando* spectroscopies, are required in order to investigate the reaction mechanisms happening on the surface of these composites under specific chemical environment and photocatalytic conditions. Nevertheless, additional theoretical and rigorous experimental studies will be helpful for the development of the next generation of both MXene- and MO-based composites with target applications in the area of photocatalysis.

Last but not least, with the focus on the abundant tunability of MXene chemical compositions, but unfortunately with a high production cost, the majority of the photocatalytic studies are still concentrating on the use of Ti-based MXenes and less studies focus on the synthesis of other MXene materials. Therefore, utilization of other MXene-based composites requires further attention in both theoretical and experimental aspects.

To finish, although MXene and MO cocatalysts are a promising and attractive alternative to replace expensive noble-metal cocatalysts in the targeted photocatalytic applications presented here, significant research efforts are still needed. Following the literature displayed in this review, it is clear that there is room for improvement especially in the design and synthesis of MXene-based composites, and in this regard, several approaches can be envisaged, such as developing more environment-friendly methods for the existing ones and extending the preparation protocols to other MXenes besides Ti-based ones.

Author contributions

M. N. and A. I. contributed to Sections 1–4, while F. N., M. F. and S. N. contributed to Sections 5 and 6, all with scientific writing, illustrations and table drafting. M. F. and S. N. supervised the research work and contributed to funding acquisition, outline drafting, reviewing, and editing the overall text, illustrations and tables of the manuscript.

Conflicts of interest

There are no conflicts to declare.



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References

- N. S. Lewis, *Science*, 2016, **351**, aad1920.
- R. Schächli, D. Rutz, F. Dähler, A. Muroyama, P. Haueter, J. Lilliestam, A. Patt, P. Furler and A. Steinfeld, *Nature*, 2022, **601**, 63–68.
- Q. Wang, C. Pornrungrroj, S. Linley and E. Reisner, *Nat. Energy*, 2022, **7**, 13–24.
- R. Wang, T. Huang, J. Xue, J. Tong, K. Zhu and Y. Yang, *Nat. Photonics*, 2021, **15**, 411–425.
- I. Massiot, A. Cattoni and S. Collin, *Nat. Energy*, 2020, **5**, 959–972.
- Y. Tachibana, L. Vayssieres and J. R. Durrant, *Nat. Photonics*, 2012, **6**, 511–518.
- J. H. Montoya, L. C. Seitz, P. Chakthranont, A. Vojvodic, T. F. Jaramillo and J. K. Nørskov, *Nat. Mater.*, 2017, **16**, 70–81.
- B. Qiu, M. Du, Y. Ma, Q. Zhu, M. Xing and J. Zhang, *Energy Environ. Sci.*, 2021, **14**, 5260–5288.
- Z. Wang, Y. Hu, S. Zhang and Y. Sun, *Chem. Soc. Rev.*, 2022, **51**, 6704–6737.
- G. Chen, G. I. N. Waterhouse, R. Shi, J. Zhao, Z. Li, L. Wu, C. Tung and T. Zhang, *Angew. Chem., Int. Ed.*, 2019, **58**, 17528–17551.
- D. G. Nocera, *Acc. Chem. Res.*, 2012, **45**, 767–776.
- R. M. Navarro Yerga, M. C. Álvarez Galván, F. del Valle, J. A. Villoria de la Mano and J. L. G. Fierro, *ChemSusChem*, 2009, **2**, 471–485.
- A. Kubacka, M. Fernández-García and G. Colón, *Chem. Rev.*, 2012, **112**, 1555–1614.
- X. Sun, S. Jiang, H. Huang, H. Li, B. Jia and T. Ma, *Angew. Chem., Int. Ed.*, 2022, **61**, e2022048.
- X. Li, J. Yu and M. Jaroniec, *Chem. Soc. Rev.*, 2016, **45**, 2603–2636.
- Z. Wang, C. Li and K. Domen, *Chem. Soc. Rev.*, 2019, **48**, 2109–2125.
- L. Jing, W. Zhou, G. Tian and H. Fu, *Chem. Soc. Rev.*, 2013, **42**, 9509.
- M. Ahmed and G. Xinxin, *Inorg. Chem. Front.*, 2016, **3**, 578–590.
- M. Z. Rahman, M. G. Kibria and C. B. Mullins, *Chem. Soc. Rev.*, 2020, **49**, 1887–1931.
- J. Yoon, J. Kim, C. Kim, H. W. Jang and J. Lee, *Adv. Energy Mater.*, 2021, **11**, 2003052.
- Q. Xiang, B. Cheng and J. Yu, *Angew. Chem., Int. Ed.*, 2015, **54**, 11350–11366.
- S. Bai, W. Yin, L. Wang, Z. Li and Y. Xiong, *RSC Adv.*, 2016, **6**, 57446–57463.
- J. Ran, M. Jaroniec and S. Qiao, *Adv. Mater.*, 2018, **30**, 1704649.
- G. Zhao and X. Xu, *Nanoscale*, 2021, **13**, 10649–10667.
- H. Zhao, L. Jian, M. Gong, M. Jing, H. Li, Q. Mao, T. Lu, Y. Guo, R. Ji, W. Chi, Y. Dong and Y. Zhu, *Small Struct.*, 2022, **3**, 2100229.
- D. Wang and X.-Q. Gong, *Nat. Commun.*, 2021, **12**, 158.
- R. Reichert, Z. Jusys and R. J. Behm, *J. Phys. Chem. C*, 2015, **119**, 24750–24759.
- X. Wang, Y. Ruan, S. Feng, S. Chen and K. Su, *ACS Sustainable Chem. Eng.*, 2018, **6**, 11424–11432.
- S. Bai, L. Yang, C. Wang, Y. Lin, J. Lu, J. Jiang and Y. Xiong, *Angew. Chem., Int. Ed.*, 2015, **54**, 14810–14814.
- H. Lin, B. Sun, H. Wang, Q. Ruan, Y. Geng, Y. Li, J. Wu, W. Wang, J. Liu and X. Wang, *Small*, 2019, **15**, 1804115.
- T. Simon, N. Bouchonville, M. J. Berr, A. Vaneski, A. Adrović, D. Volbers, R. Wyrwich, M. Döblinger, A. S. Susha, A. L. Rogach, F. Jäckel, J. K. Stolarczyk and J. Feldmann, *Nat. Mater.*, 2014, **13**, 1013–1018.
- G. Ge, M. Liu, C. Liu, W. Zhou, D. Wang, L. Liu and J. Ye, *J. Mater. Chem. A*, 2019, **7**, 9222–9229.
- Z. Sun, H. Zheng, J. Li and P. Du, *Energy Environ. Sci.*, 2015, **8**, 2668–2676.
- R. Tang, S. Xiong, D. Gong, Y. Deng, Y. Wang, L. Su, C. Ding, L. Yang and C. Liao, *ACS Appl. Mater. Interfaces*, 2020, **12**, 56663–56680.
- L. Cheng, X. Li, H. Zhang and Q. Xiang, *J. Phys. Chem. Lett.*, 2019, **10**, 3488–3494.
- Y. Zhao, M. Que, J. Chen and C. Yang, *J. Mater. Chem. C*, 2020, **8**, 16258–16281.
- X. Xie and N. Zhang, *Adv. Funct. Mater.*, 2020, **30**, 2002528.
- K. Huang, C. Li, H. Li, G. Ren, L. Wang, W. Wang and X. Meng, *ACS Appl. Nano Mater.*, 2020, **3**, 9581–9603.
- X. Li, Y. Bai, X. Shi, N. Su, G. Nie, R. Zhang, H. Nie and L. Ye, *Mater. Adv.*, 2021, **2**, 1570–1594.
- M. Naguib, M. W. Barsoum and Y. Gogotsi, *Adv. Mater.*, 2021, **33**, 2103393.
- K. R. G. Lim, M. Shekhirev, B. C. Wyatt, B. Anasori, Y. Gogotsi and Z. W. Seh, *Nat. Synth.*, 2022, **1**, 601–614.
- Á. Morales-García, F. Calle-Vallejo and F. Illas, *ACS Catal.*, 2020, **10**, 13487–13503.
- R. Wang, M. Li, K. Sun, Y. Zhang, J. Li and W. Bao, *Small*, 2022, **18**, 2201740.
- F. Bu, M. M. Zagho, Y. Ibrahim, B. Ma, A. Elzatahry and D. Zhao, *Nano Today*, 2020, **30**, 100803.
- I. M. Chirica, A. G. Mirea, Ş. Neaţu, M. Florea, M. W. Barsoum and F. Neaţu, *J. Mater. Chem. A*, 2021, **9**, 19589–19612.
- Y. Zhao, S. Zhang, R. Shi, G. I. N. Waterhouse, J. Tang and T. Zhang, *Mater. Today*, 2020, **34**, 78–91.
- X. Meng, N. Yun and Z. Zhang, *Can. J. Chem. Eng.*, 2019, **97**, 1982–1998.
- A. K. Singh, K. Mathew, H. L. Zhuang and R. G. Hennig, *J. Phys. Chem. Lett.*, 2015, **6**, 1087–1098.
- X. Zhou and H. Dong, *ChemCatChem*, 2019, **11**, 3688–3715.



- 50 P. Liao and E. A. Carter, *Chem. Soc. Rev.*, 2013, **42**, 2401–2422.
- 51 C. Zhang, G. Chen, Y. Si and M. Liu, *Phys. Chem. Chem. Phys.*, 2022, **24**, 1237–1261.
- 52 Y. Dong, P. Duchesne, A. Mohan, K. K. Ghuman, P. Kant, L. Hurtado, U. Ulmer, J. Y. Y. Loh, A. A. Tountas, L. Wang, A. Jelle, M. Xia, R. Dittmeyer and G. A. Ozin, *Chem. Soc. Rev.*, 2020, **49**, 5648–5663.
- 53 H. Tong, S. Ouyang, Y. Bi, N. Umezawa, M. Oshikiri and J. Ye, *Adv. Mater.*, 2012, **24**, 229–251.
- 54 M. Humayun, C. Wang and W. Luo, *Small Methods*, 2022, **6**, 2101395.
- 55 D. M. Alonso, S. G. Wettstein and J. A. Dumesic, *Chem. Soc. Rev.*, 2012, **41**, 8075–8098.
- 56 R. Huang, X. Li, W. Gao, X. Zhang, S. Liang and M. Luo, *RSC Adv.*, 2021, **11**, 14844–14861.
- 57 L. Luo, T. Zhang, M. Wang, R. Yun and X. Xiang, *ChemSusChem*, 2020, **13**, 5173–5184.
- 58 C. C. Nguyen, N. N. Vu and T.-O. Do, *J. Mater. Chem. A*, 2015, **3**, 18345–18359.
- 59 M. Xiao, Z. Wang, M. Lyu, B. Luo, S. Wang, G. Liu, H. Cheng and L. Wang, *Adv. Mater.*, 2019, **31**, 1801369.
- 60 D. Chen, X. Zhang and A. F. Lee, *J. Mater. Chem. A*, 2015, **3**, 14487–14516.
- 61 H. Li, Z. Bian, J. Zhu, D. Zhang, G. Li, Y. Huo, H. Li and Y. Lu, *J. Am. Chem. Soc.*, 2007, **129**, 8406–8407.
- 62 X. Wang, M. Liao, Y. Zhong, J. Y. Zheng, W. Tian, T. Zhai, C. Zhi, Y. Ma, J. Yao, Y. Bando and D. Golberg, *Adv. Mater.*, 2012, **24**, 3421–3425.
- 63 T. Xiong, F. Dong and Z. Wu, *RSC Adv.*, 2014, **4**, 56307–56312.
- 64 G. Prieto, H. Tüysüz, N. Duyckaerts, J. Knossalla, G.-H. Wang and F. Schüth, *Chem. Rev.*, 2016, **116**, 14056–14119.
- 65 F. E. Osterloh, *Chem. Soc. Rev.*, 2013, **42**, 2294–2320.
- 66 M. Naguib, M. Kurtoglu, V. Presser, J. Lu, J. Niu, M. Heon, L. Hultman, Y. Gogotsi and M. W. Barsoum, *Adv. Mater.*, 2011, **23**, 4248–4253.
- 67 A. VahidMohammadi, J. Rosen and Y. Gogotsi, *Science*, 2021, **372**, eabf1581.
- 68 G. Deysher, C. E. Shuck, K. Hantanasirisakul, N. C. Frey, A. C. Foucher, K. Maleski, A. Sarycheva, V. B. Shenoy, E. A. Stach, B. Anasori and Y. Gogotsi, *ACS Nano*, 2020, **14**, 204–217.
- 69 O. Mashtalir, M. Naguib, V. N. Mochalin, Y. Dall'Agnese, M. Heon, M. W. Barsoum and Y. Gogotsi, *Nat. Commun.*, 2013, **4**, 1716.
- 70 M. Naguib, R. R. Unocic, B. L. Armstrong and J. Nanda, *Dalton Trans.*, 2015, **44**, 9353–9358.
- 71 M. Ghidui, M. R. Lukatskaya, M.-Q. Zhao, Y. Gogotsi and M. W. Barsoum, *Nature*, 2014, **516**, 78–81.
- 72 J. Halim, M. R. Lukatskaya, K. M. Cook, J. Lu, C. R. Smith, L.-Å. Näslund, S. J. May, L. Hultman, Y. Gogotsi, P. Eklund and M. W. Barsoum, *Chem. Mater.*, 2014, **26**, 2374–2381.
- 73 W. Sun, S. A. Shah, Y. Chen, Z. Tan, H. Gao, T. Habib, M. Radovic and M. J. Green, *J. Mater. Chem. A*, 2017, **5**, 21663–21668.
- 74 T. Li, L. Yao, Q. Liu, J. Gu, R. Luo, J. Li, X. Yan, W. Wang, P. Liu, B. Chen, W. Zhang, W. Abbas, R. Naz and D. Zhang, *Angew. Chem., Int. Ed.*, 2018, **57**, 6115–6119.
- 75 H. Shi, P. Zhang, Z. Liu, S. W. Park, M. R. Lohe, Y. Wu, A. Shaygan Nia, S. Yang and X. Feng, *Angew. Chem., Int. Ed.*, 2021, **60**, 8689–8693.
- 76 Y. Li, H. Shao, Z. Lin, J. Lu, L. Liu, B. Duployer, P. O. Å. Persson, P. Eklund, L. Hultman, M. Li, K. Chen, X.-H. Zha, S. Du, P. Rozier, Z. Chai, E. Raymundo-Piñero, P.-L. Taberna, P. Simon and Q. Huang, *Nat. Mater.*, 2020, **19**, 894–899.
- 77 M. Li, J. Lu, K. Luo, Y. Li, K. Chang, K. Chen, J. Zhou, J. Rosen, L. Hultman, P. Eklund, P. O. Å. Persson, S. Du, Z. Chai, Z. Huang and Q. Huang, *J. Am. Chem. Soc.*, 2019, **141**, 4730–4737.
- 78 B. Anasori, Y. Xie, M. Beidaghi, J. Lu, B. C. Hosler, L. Hultman, P. R. C. Kent, Y. Gogotsi and M. W. Barsoum, *ACS Nano*, 2015, **9**, 9507–9516.
- 79 Q. Tao, M. Dahlgqvist, J. Lu, S. Kota, R. Meshkian, J. Halim, J. Palisaitis, L. Hultman, M. W. Barsoum, P. O. Å. Persson and J. Rosen, *Nat. Commun.*, 2017, **8**, 14949.
- 80 L. Dong, H. Kumar, B. Anasori, Y. Gogotsi and V. B. Shenoy, *J. Phys. Chem. Lett.*, 2017, **8**, 422–428.
- 81 N. M. Caffrey, *J. Phys. Chem. C*, 2020, **124**, 18797–18804.
- 82 J. Halim, A. S. Etman, A. Elsukova, P. Polcik, J. Palisaitis, M. W. Barsoum, P. O. Å. Persson and J. Rosen, *Nanoscale*, 2021, **13**, 311–319.
- 83 R. Meshkian, H. Lind, J. Halim, A. El Ghazaly, J. Thörnberg, Q. Tao, M. Dahlgqvist, J. Palisaitis, P. O. Å. Persson and J. Rosen, *ACS Appl. Nano Mater.*, 2019, **2**, 6209–6219.
- 84 M. Zhao, X. Xie, C. E. Ren, T. Makaryan, B. Anasori, G. Wang and Y. Gogotsi, *Adv. Mater.*, 2017, **29**, 1702410.
- 85 C. E. Ren, M. Zhao, T. Makaryan, J. Halim, M. Boota, S. Kota, B. Anasori, M. W. Barsoum and Y. Gogotsi, *ChemElectroChem*, 2016, **3**, 689–693.
- 86 M. R. Lukatskaya, S. Kota, Z. Lin, M.-Q. Zhao, N. Shpigel, M. D. Levi, J. Halim, P.-L. Taberna, M. W. Barsoum, P. Simon and Y. Gogotsi, *Nat. Energy*, 2017, **2**, 17105.
- 87 L. Xiu, Z. Wang, M. Yu, X. Wu and J. Qiu, *ACS Nano*, 2018, **12**, 8017–8028.
- 88 Y. Xia, T. S. Mathis, M.-Q. Zhao, B. Anasori, A. Dang, Z. Zhou, H. Cho, Y. Gogotsi and S. Yang, *Nature*, 2018, **557**, 409–412.
- 89 X. Xie, K. Kretschmer, B. Anasori, B. Sun, G. Wang and Y. Gogotsi, *ACS Appl. Nano Mater.*, 2018, **1**, 505–511.
- 90 Q. Zhao, Q. Zhu, J. Miao, P. Zhang, P. Wan, L. He and B. Xu, *Small*, 2019, **15**, 1904293.
- 91 Q. Xue, H. Zhang, M. Zhu, Z. Pei, H. Li, Z. Wang, Y. Huang, Y. Huang, Q. Deng, J. Zhou, S. Du, Q. Huang and C. Zhi, *Adv. Mater.*, 2017, **29**, 1604847.
- 92 Q. Zhang, Y. Sun, M. Liu and Y. Liu, *Nanoscale*, 2020, **12**, 1826–1832.
- 93 B. Shao, Z. Liu, G. Zeng, H. Wang, Q. Liang, Q. He, M. Cheng, C. Zhou, L. Jiang and B. Song, *J. Mater. Chem. A*, 2020, **8**, 7508–7535.



- 94 H. Mao, C. Gu, S. Yan, Q. Xin, S. Cheng, P. Tan, X. Wang, F. Xiu, X. Liu, J. Liu, W. Huang and L. Sun, *Adv. Electron. Mater.*, 2020, **6**, 1900493.
- 95 T. Su, R. Peng, Z. D. Hood, M. Naguib, I. N. Ivanov, J. K. Keum, Z. Qin, Z. Guo and Z. Wu, *ChemSusChem*, 2018, **11**, 688–699.
- 96 Y. Li, D. Zhang, X. Feng, Y. Liao, Q. Wen and Q. Xiang, *Nanoscale Adv.*, 2019, **1**, 1812–1818.
- 97 H. Wang, R. Peng, Z. D. Hood, M. Naguib, S. P. Adhikari and Z. Wu, *ChemSusChem*, 2016, **9**, 1490–1497.
- 98 T. Su, Z. D. Hood, M. Naguib, L. Bai, S. Luo, C. M. Rouleau, I. N. Ivanov, H. Ji, Z. Qin and Z. Wu, *ACS Appl. Energy Mater.*, 2019, **2**, 4640–4651.
- 99 T. Su, Z. D. Hood, M. Naguib, L. Bai, S. Luo, C. M. Rouleau, I. N. Ivanov, H. Ji, Z. Qin and Z. Wu, *Nanoscale*, 2019, **11**, 8138–8149.
- 100 K. Maeda, H. Wakayama, Y. Washio, A. Ishikawa, M. Okazaki, H. Nakata and S. Matsuishi, *J. Phys. Chem. C*, 2020, **124**, 14640–14645.
- 101 Z. Guo, J. Zhou, L. Zhu and Z. Sun, *J. Mater. Chem. A*, 2016, **4**, 11446–11452.
- 102 X. Zhang, X. Zhao, D. Wu, Y. Jing and Z. Zhou, *Nanoscale*, 2015, **7**, 16020–16025.
- 103 G. Gao, A. P. O'Mullane and A. Du, *ACS Catal.*, 2017, **7**, 494–500.
- 104 K. Xiong, P. Wang, G. Yang, Z. Liu, H. Zhang, S. Jin and X. Xu, *Sci. Rep.*, 2017, **7**, 15095.
- 105 G. Jia, Y. Wang, X. Cui and W. Zheng, *ACS Sustainable Chem. Eng.*, 2018, **6**, 13480–13486.
- 106 A. Fujishima and K. Honda, *Nature*, 1972, **238**, 37–38.
- 107 T. Inoue, A. Fujishima, S. Konishi and K. Honda, *Nature*, 1979, **277**, 637–638.
- 108 Y. Li, X. Deng, J. Tian, Z. Liang and H. Cui, *Appl. Mater. Today*, 2018, **13**, 217–227.
- 109 C. Peng, X. Xie, W. Xu, T. Zhou, P. Wei, J. Jia, K. Zhang, Y. Cao, H. Wang, F. Peng, R. Yang, X. Yan, H. Pan and H. Yu, *Chem. Eng. J.*, 2021, **421**, 128766.
- 110 B. Wang, M. Wang, F. Liu, Q. Zhang, S. Yao, X. Liu and F. Huang, *Angew. Chem., Int. Ed.*, 2020, **59**, 1914–1918.
- 111 J. Ran, G. Gao, F.-T. Li, T.-Y. Ma, A. Du and S.-Z. Qiao, *Nat. Commun.*, 2017, **8**, 13907.
- 112 R. Xiao, C. Zhao, Z. Zou, Z. Chen, L. Tian, H. Xu, H. Tang, Q. Liu, Z. Lin and X. Yang, *Appl. Catal., B*, 2020, **268**, 118382.
- 113 B. Sun, P. Qiu, Z. Liang, Y. Xue, X. Zhang, L. Yang, H. Cui and J. Tian, *Chem. Eng. J.*, 2021, **406**, 127177.
- 114 M. Ding, R. Xiao, C. Zhao, D. Bukhvalov, Z. Chen, H. Xu, H. Tang, J. Xu and X. Yang, *Sol. RRL*, 2021, **5**, 2000414.
- 115 N. Zhao, Y. Hu, J. Du, G. Liu, B. Dong, Y. Yang, J. Peng, J. Li and M. Zhai, *Appl. Surf. Sci.*, 2020, **530**, 147247.
- 116 S. Jin, Z. Shi, H. Jing, L. Wang, Q. Hu, D. Chen, N. Li and A. Zhou, *ACS Appl. Energy Mater.*, 2021, **4**, 12754–12766.
- 117 G. Zuo, Y. Wang, W. L. Teo, A. Xie, Y. Guo, Y. Dai, W. Zhou, D. Jana, Q. Xian, W. Dong and Y. Zhao, *Angew. Chem., Int. Ed.*, 2020, **59**, 11287–11292.
- 118 X. Du, T. Zhao, Z. Xiu, Z. Xing, Z. Li, K. Pan, S. Yang and W. Zhou, *Appl. Mater. Today*, 2020, **20**, 100719.
- 119 K. Huang, C. Li and X. Meng, *J. Colloid Interface Sci.*, 2020, **580**, 669–680.
- 120 R. Chen, P. Wang, J. Chen, C. Wang and Y. Ao, *Appl. Surf. Sci.*, 2019, **473**, 11–19.
- 121 Y. Li, L. Ding, Z. Liang, Y. Xue, H. Cui and J. Tian, *Chem. Eng. J.*, 2020, **383**, 123178.
- 122 Z. Li, W. Huang, J. Liu, K. Lv and Q. Li, *ACS Catal.*, 2021, **11**, 8510–8520.
- 123 Y. Li, L. Ding, Y. Guo, Z. Liang, H. Cui and J. Tian, *ACS Appl. Mater. Interfaces*, 2019, **11**, 41440–41447.
- 124 J. Kang, S. Byun, S. Kim, J. Lee, M. Jung, H. Hwang, T. W. Kim, S. H. Song and D. Lee, *ACS Appl. Energy Mater.*, 2020, **3**, 9226–9233.
- 125 J. Li, L. Zhao, S. Wang, J. Li, G. Wang and J. Wang, *Appl. Surf. Sci.*, 2020, **515**, 145922.
- 126 T. Song, L. Hou, B. Long, A. Ali and G. J. Deng, *J. Colloid Interface Sci.*, 2021, **584**, 474–483.
- 127 Y. Li, X. Chen, Y. Sun, X. Meng, Y. Dall'Agnese, G. Chen, C. Dall'Agnese, H. Ren, S. Sasaki, H. Tamiaki and X. Wang, *Adv. Mater. Interfaces*, 2020, **7**, 1902080.
- 128 Y. Li, Y. Sun, T. Zheng, Y. Dall'Agnese, C. Dall'Agnese, X. Meng, S. Sasaki, H. Tamiaki and X. Wang, *Chem. – Eur. J.*, 2021, **27**, 5277–5282.
- 129 S. Debow, T. Zhang, X. Liu, F. Song, Y. Qian, J. Han, K. Maleski, Z. B. Zander, W. R. Creasy, D. L. Kuhn, Y. Gogotsi, B. G. DeLacy and Y. Rao, *J. Phys. Chem. C*, 2021, **125**, 10473–10482.
- 130 J. Low, L. Zhang, T. Tong, B. Shen and J. Yu, *J. Catal.*, 2018, **361**, 255–266.
- 131 M. Ye, X. Wang, E. Liu, J. Ye and D. Wang, *ChemSusChem*, 2018, **11**, 1606–1611.
- 132 Q. Song, B. Shen, J. Yu and S. Cao, *ChemNanoMat*, 2021, **7**, 910–915.
- 133 Q. Tang, P. Xiong, H. Wang and Z. Wu, *J. Colloid Interface Sci.*, 2022, **619**, 179–187.
- 134 K. Wang, X. Li, N. Wang, Q. Shen, M. Liu, J. Zhou and N. Li, *Ind. Eng. Chem. Res.*, 2021, **60**, 8720–8732.
- 135 Z. Zeng, Y. Yan, J. Chen, P. Zan, Q. Tian and P. Chen, *Adv. Funct. Mater.*, 2019, **29**, 1806500.
- 136 J. Zhang, J. Shi, S. Tao, L. Wu and J. Lu, *Appl. Surf. Sci.*, 2021, **542**, 148685.
- 137 S. Cao, B. Shen, T. Tong, J. Fu and J. Yu, *Adv. Funct. Mater.*, 2018, **28**, 1800136.
- 138 J. Li, Z. Wang, H. Chen, Q. Zhang, H. Hu, L. Liu, J. Ye and D. Wang, *Catal. Sci. Technol.*, 2021, **11**, 4953–4961.
- 139 C. Yang, Q. Tan, Q. Li, J. Zhou, J. Fan, B. Li, J. Sun and K. Lv, *Appl. Catal., B*, 2020, **268**, 118738.
- 140 F. He, B. Zhu, B. Cheng, J. Yu, W. Ho and W. Macyk, *Appl. Catal., B*, 2020, **272**, 119006.
- 141 Q. Tang, Z. Sun, S. Deng, H. Wang and Z. Wu, *J. Colloid Interface Sci.*, 2020, **564**, 406–417.
- 142 H. Wang, Q. Tang and Z. Wu, *ACS Sustainable Chem. Eng.*, 2021, **9**, 8425–8434.
- 143 L. Hong, R. Guo, Y. Yuan, X. Ji, Z. Lin, X. Yin and W. Pan, *Colloids Surf., A*, 2022, **639**, 128358.
- 144 W. Chen, B. Han, Y. Xie, S. Liang, H. Deng and Z. Lin, *Chem. Eng. J.*, 2020, **391**, 123519.



- 145 Q. Shi, X. Zhang, Y. Yang, J. Huang, X. Fu, T. Wang, X. Liu, A. Sun, J. Ge, J. Shen, Y. Zhou and Z. Liu, *J. Energy Chem.*, 2021, **59**, 9–18.
- 146 M. Que, Y. Zhao, Y. Yang, L. Pan, W. Lei, W. Cai, H. Yuan, J. Chen and G. Zhu, *ACS Appl. Mater. Interfaces*, 2021, **13**, 6180–6187.
- 147 M. Que, W. Cai, Y. Zhao, Y. Yang, B. Zhang, S. Yun, J. Chen and G. Zhu, *J. Colloid Interface Sci.*, 2022, **610**, 538–545.
- 148 Z. Zhang, B. Wang, H.-B. Zhao, J.-F. Liao, Z.-C. Zhou, T. Liu, B. He, Q. Wei, S. Chen, H.-Y. Chen, D.-B. Kuang, Y. Li and G. Xing, *Appl. Catal., B*, 2022, **312**, 121358.
- 149 H. Liu, *Chin. J. Catal.*, 2014, **35**, 1619–1640.
- 150 S. D. Minter, P. Christopher and S. Linic, *ACS Energy Lett.*, 2019, **4**, 163–166.
- 151 A. Liu, Y. Yang, X. Ren, Q. Zhao, M. Gao, W. Guan, F. Meng, L. Gao, Q. Yang, T. Ma and X. Liang, *ChemSusChem*, 2020, **13**, 3766–3788.
- 152 J. A. Pool, E. Lobkovsky and P. J. Chirik, *Nature*, 2004, **427**, 527–530.
- 153 J. Qin, X. Hu, X. Li, Z. Yin, B. Liu and K. Lam, *Nano Energy*, 2019, **61**, 27–35.
- 154 B. Chang, Y. Guo, D. Wu, L. Li, B. Yang and J. Wang, *Chem. Sci.*, 2021, **12**, 11213–11224.
- 155 C. Hao, Y. Liao, Y. Wu, Y. An, J. Lin, Z. Gu, M. Jiang, S. Hu and X. Wang, *J. Phys. Chem. Solids*, 2020, **136**, 109141.
- 156 W. Gao, X. Li, S. Luo, Z. Luo, X. Zhang, R. Huang and M. Luo, *J. Colloid Interface Sci.*, 2021, **585**, 20–29.
- 157 H. Jiang, C. Zang, Y. Zhang, W. Wang, C. Yang, B. Sun, Y. Shen and F. Bian, *Catal. Sci. Technol.*, 2020, **10**, 5964–5972.
- 158 Q. Liu, L. Ai and J. Jiang, *J. Mater. Chem. A*, 2018, **6**, 4102–4110.
- 159 J. Qian, S. Zhao, W. Dang, Y. Liao, W. Zhang, H. Wang, L. Lv, L. Luo, H.-Y. Jiang and J. Tang, *Adv. Sustainable Syst.*, 2021, **5**, 2000282.
- 160 T. Hou, Q. Li, Y. Zhang, W. Zhu, K. Yu, S. Wang, Q. Xu, S. Liang and L. Wang, *Appl. Catal., B*, 2020, **273**, 119072.
- 161 L. I. Qi, J. Xu, S. Wang, K. Yu, W. Zhu, T. Hou, L. Zhang, W. Zhang, S. Liang and L. Wang, *Trans. Nonferrous Met. Soc. China*, 2022, **32**, 233–250.
- 162 B. Chang, Y. Guo, H. Liu, L. Li and B. Yang, *J. Mater. Chem. A*, 2022, **10**, 3134–3145.
- 163 H. Li, J. Shang, Z. Ai and L. Zhang, *J. Am. Chem. Soc.*, 2015, **137**, 6393–6399.
- 164 W. Song, E. M. Lopato, S. Bernhard, P. A. Salvador and G. S. Rohrer, *Appl. Catal., B*, 2020, **269**, 118750.
- 165 S. Martha, P. Chandra Sahoo and K. M. Parida, *RSC Adv.*, 2015, **5**, 61535–61553.
- 166 Y. Sun, T. Zhang, C. Li, K. Xu and Y. Li, *J. Mater. Chem. A*, 2020, **8**, 13415–13436.
- 167 Y. Yang, S. Niu, D. Han, T. Liu, G. Wang and Y. Li, *Adv. Energy Mater.*, 2017, **7**, 1700555.
- 168 X. Li, H. Zhao, J. Liang, Y. Luo, G. Chen, X. Shi, S. Lu, S. Gao, J. Hu, Q. Liu and X. Sun, *J. Mater. Chem. A*, 2021, **9**, 6650–6670.
- 169 Z. Zeng, Y. Xu, Z. Zhang, Z. Gao, M. Luo, Z. Yin, C. Zhang, J. Xu, B. Huang, F. Luo, Y. Du and C. Yan, *Chem. Soc. Rev.*, 2020, **49**, 1109–1143.
- 170 M. A. Peña and J. L. G. Fierro, *Chem. Rev.*, 2001, **101**, 1981–2018.
- 171 S. Royer, D. Duprez, F. Can, X. Courtois, C. Batiot-Dupeyrat, S. Laassiri and H. Alamdari, *Chem. Rev.*, 2014, **114**, 10292–10368.
- 172 A. E. Danks, S. R. Hall and Z. Schnepf, *Mater. Horiz.*, 2016, **3**, 91–112.
- 173 D. R. Modeshia and R. I. Walton, *Chem. Soc. Rev.*, 2010, **39**, 4303.
- 174 X. Huang, G. Zhao, G. Wang and J. T. S. Irvine, *Chem. Sci.*, 2018, **9**, 3623–3637.
- 175 Y. Jun, J. Choi and J. Cheon, *Angew. Chem., Int. Ed.*, 2006, **45**, 3414–3439.
- 176 Y. Mao, T.-J. Park, F. Zhang, H. Zhou and S. S. Wong, *Small*, 2007, **3**, 1122–1139.
- 177 M.-M. Titirici, M. Antonietti and A. Thomas, *Chem. Mater.*, 2006, **18**, 3808–3812.
- 178 A. Querejeta, A. Varela, M. Parras, F. del Monte, M. García-Hernández and J. M. González-Calbet, *Chem. Mater.*, 2009, **21**, 1898–1905.
- 179 S. Bie, Y. Zhu, J. Su, C. Jin, S. Liu, R. Yang and J. Wu, *J. Mater. Chem. A*, 2015, **3**, 22448–22453.
- 180 J. J. Urban, L. Ouyang, M.-H. Jo, D. S. Wang and H. Park, *Nano Lett.*, 2004, **4**, 1547–1550.
- 181 C. T. Kresge, M. E. Leonowicz, W. J. Roth, J. C. Vartuli and J. S. Beck, *Nature*, 1992, **359**, 710–712.
- 182 D. Gu and F. Schüth, *Chem. Soc. Rev.*, 2014, **43**, 313–344.
- 183 H. Yang and D. Zhao, *J. Mater. Chem.*, 2005, **15**, 1217–1231.
- 184 A.-H. Lu and F. Schüth, *Adv. Mater.*, 2006, **18**, 1793–1805.
- 185 F. Schüth, *Angew. Chem., Int. Ed.*, 2003, **42**, 3604–3622.
- 186 M. M. Nair, H. Yen and F. Kleitz, *C. R. Chim.*, 2014, **17**, 641–655.
- 187 M. M. Nair, S. Kaliaguine and F. Kleitz, *ACS Catal.*, 2014, **4**, 3837–3846.
- 188 Y. Shi, Y. Wan and D. Zhao, *Chem. Soc. Rev.*, 2011, **40**, 3854.
- 189 B. T. Yonemoto, G. S. Hutchings and F. Jiao, *J. Am. Chem. Soc.*, 2014, **136**, 8895–8898.
- 190 J. Liu, Q. Jia, J. Long, X. Wang, Z. Gao and Q. Gu, *Appl. Catal., B*, 2018, **222**, 35–43.
- 191 G. D. Moon, J. B. Joo, I. Lee and Y. Yin, *Nanoscale*, 2014, **6**, 12002–12008.
- 192 P. Ravi, V. Navakoteswara Rao, M. V. Shankar and M. Sathish, *Int. J. Hydrogen Energy*, 2020, **45**, 7517–7529.
- 193 X. Chen, W. Chen, H. Gao, Y. Yang and W. Shangguan, *Appl. Catal., B*, 2014, **152–153**, 68–72.
- 194 B. Fang, Y. Xing, A. Bonakdarpour, S. Zhang and D. P. Wilkinson, *ACS Sustainable Chem. Eng.*, 2015, **3**, 2381–2388.
- 195 M. M. Gui, S.-P. Chai and A. R. Mohamed, *Appl. Surf. Sci.*, 2014, **319**, 37–43.
- 196 X. Chen, W. Chen, P. Lin, Y. Yang, H. Gao, J. Yuan and W. Shangguan, *Catal. Commun.*, 2013, **36**, 104–108.
- 197 X. Ning, W. Zhen, X. Zhang and G. Lu, *ChemSusChem*, 2019, **12**, 1410–1420.



- 198 Y. Fu, C. Liu, C. Zhu, H. Wang, Y. Dou, W. Shi, M. Shao, H. Huang, Y. Liu and Z. Kang, *Inorg. Chem. Front.*, 2018, **5**, 1646–1652.
- 199 N. Banic, J. Krstić, S. Stojadinović, A. Brnović, A. Djordjevic and B. Abramović, *Int. J. Energy Res.*, 2020, **44**, 8951–8963.
- 200 L. Li, B. Cheng, Y. Wang and J. Yu, *J. Colloid Interface Sci.*, 2015, **449**, 115–121.
- 201 N. Chen, J. Cao, M. Guo, C. Liu, H. Lin and S. Chen, *Int. J. Hydrogen Energy*, 2021, **46**, 19363–19372.
- 202 B. Yan, J. Li, Z. Lin, C. Du and G. Yang, *ACS Appl. Nano Mater.*, 2019, **2**, 6783–6792.
- 203 J. Chen, S. Shen, P. Guo, M. Wang, P. Wu, X. Wang and L. Guo, *Appl. Catal., B*, 2014, **152–153**, 335–341.
- 204 L. Zhu, M. Hong and G. W. Ho, *Nano Energy*, 2015, **11**, 28–37.
- 205 G. Sadanandam, X. Luo, X. Chen, Y. Bao, K. P. Homewood and Y. Gao, *Appl. Surf. Sci.*, 2021, **541**, 148687.
- 206 Z. Mao, J. Chen, Y. Yang, D. Wang, L. Bie and B. D. Fahlman, *ACS Appl. Mater. Interfaces*, 2017, **9**, 12427–12435.
- 207 Y. Zhu, T. Wan, X. Wen, D. Chu and Y. Jiang, *Appl. Catal., B*, 2019, **244**, 814–822.
- 208 E. L. Uzunova, N. Seriani and H. Mikosch, *Phys. Chem. Chem. Phys.*, 2015, **17**, 11088–11094.
- 209 S. Zhu, S. Liang, Y. Tong, X. An, J. Long, X. Fu and X. Wang, *Phys. Chem. Chem. Phys.*, 2015, **17**, 9761–9770.
- 210 H. Li, Y. Lei, Y. Huang, Y. Fang, Y. Xu, L. Zhu and X. Li, *J. Nat. Gas Chem.*, 2011, **20**, 145–150.
- 211 Y. Li, W. Zhang, X. Shen, P. Peng, L. Xiong and Y. Yu, *Chin. J. Catal.*, 2015, **36**, 2229–2236.
- 212 M. Tahir and B. Tahir, *Appl. Surf. Sci.*, 2016, **377**, 244–252.
- 213 S. Shoji, A. Yamaguchi, E. Sakai and M. Miyauchi, *ACS Appl. Mater. Interfaces*, 2017, **9**, 20613–20619.
- 214 M. Alves Melo Júnior, A. Morais and A. F. Nogueira, *Microporous Mesoporous Mater.*, 2016, **234**, 1–11.
- 215 V. Jeyalakshmi, R. Mahalakshmy, K. R. Krishnamurthy and B. Viswanathan, *Catal. Today*, 2016, **266**, 160–167.
- 216 L. F. Garay-Rodríguez and L. M. Torres-Martínez, *J. Mater. Sci.: Mater. Electron.*, 2020, **31**, 19248–19265.
- 217 S. Jeong, W. D. Kim, S. Lee, K. Lee, S. Lee, D. Lee and D. C. Lee, *ChemCatChem*, 2016, **8**, 1641–1645.
- 218 Z.-Y. Wang, H.-C. Chou, J. C. S. Wu, D. P. Tsai and G. Mul, *Appl. Catal., A*, 2010, **380**, 172–177.
- 219 D.-S. Lee, H.-J. Chen and Y.-W. Chen, *J. Phys. Chem. Solids*, 2012, **73**, 661–669.
- 220 X. Shao, X. Yin and J. Wang, *J. Colloid Interface Sci.*, 2018, **512**, 466–473.
- 221 D.-S. Lee and Y.-W. Chen, *J. CO2 Util.*, 2015, **10**, 1–6.
- 222 C.-W. Tsai, H. M. Chen, R.-S. Liu, K. Asakura and T.-S. Chan, *J. Phys. Chem. C*, 2011, **115**, 10180–10186.
- 223 F. Wang, Y. Zhou, P. Li, L. Kuai and Z. Zou, *Chin. J. Catal.*, 2016, **37**, 863–868.
- 224 J. A. Torres, A. E. Nogueira, G. T. S. T. da Silva, O. F. Lopes, Y. Wang, T. He and C. Ribeiro, *J. CO2 Util.*, 2020, **35**, 106–114.
- 225 S. W. Jo, B. S. Kwak, K. M. Kim, J. Y. Do, N.-K. Park, S. O. Ryu, H.-J. Ryu, J.-I. Baek and M. Kang, *Appl. Surf. Sci.*, 2015, **355**, 891–901.
- 226 X. Xue, R. Chen, C. Yan, Y. Hu, W. Zhang, S. Yang, L. Ma, G. Zhu and Z. Jin, *Nanoscale*, 2019, **11**, 10439–10445.
- 227 B. Sun, Z. Liang, Y. Qian, X. Xu, Y. Han and J. Tian, *ACS Appl. Mater. Interfaces*, 2020, **12**, 7257–7269.
- 228 X. Hui, L. Li, Q. Xia, S. Hong, L. Hao, A. W. Robertson and Z. Sun, *Chem. Eng. J.*, 2022, **438**, 135485.
- 229 H. Mou, J. Wang, D. Zhang, D. Yu, W. Chen, D. Wang and T. Mu, *J. Mater. Chem. A*, 2019, **7**, 5719–5725.
- 230 L. Yan, Q. Zhang, W. Deng, Q. Zhang and Y. Wang, *Catalytic Valorization of Biomass and Bioplatfoms to Chemicals through Deoxygenation*, Elsevier Inc., 1st edn, 2020, vol. 66.
- 231 H. H. B. T.-S, in *Transition Metal Oxides*, ed. S. S. and C. Kung, Elsevier, 1989, vol. 45, pp. 121–135.
- 232 M. Naguib, M. Kurtoglu, V. Presser, J. Lu, J. Niu, M. Heon, L. Hultman, Y. Gogotsi and M. W. Barsoum, *Adv. Mater.*, 2011, **23**, 4248.
- 233 M. Khazaei, M. Arai, T. Sasaki, A. Ranjbar, Y. Liang and S. Yunoki, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2015, **92**, 75411.
- 234 C. Peng, X. Yang, Y. Li, H. Yu, H. Wang and F. Peng, *ACS Appl. Mater. Interfaces*, 2016, **8**, 6051–6060.
- 235 C. Peng, H. Wang, H. Yu and F. Peng, *Mater. Res. Bull.*, 2017, **89**, 16–25.
- 236 Y. Liu, H. Xiao and W. A. Goddard, *J. Am. Chem. Soc.*, 2016, **138**, 15853–15856.
- 237 C. Peng, P. Wei, X. Li, Y. Liu, Y. Cao, H. Wang, H. Yu, F. Peng, L. Zhang, B. Zhang and K. Lv, *Nano Energy*, 2018, **53**, 97–107.
- 238 J. K. Nørskov, T. Bligaard, J. Rossmeisl and C. H. Christensen, *Nat. Chem.*, 2009, **1**, 37–46.
- 239 B. Hinnemann, P. G. Moses, J. Bonde, K. P. Jørgensen, J. H. Nielsen, S. Horch, I. Chorkendorff and J. K. Nørskov, *J. Am. Chem. Soc.*, 2005, **127**, 5308–5309.
- 240 J. Bonde, P. G. Moses, T. F. Jaramillo, J. K. Nørskov and I. Chorkendorff, *Faraday Discuss.*, 2009, **140**, 219–231.
- 241 Y. H. Li, C. Peng, S. Yang, H. F. Wang and H. G. Yang, *J. Catal.*, 2015, **330**, 120–128.
- 242 B. Weng, M.-Y. Qi, C. Han, Z.-R. Tang and Y.-J. Xu, *ACS Catal.*, 2019, **9**, 4642–4687.
- 243 T. Cai, L. Wang, Y. Liu, S. Zhang, W. Dong, H. Chen, X. Yi, J. Yuan, X. Xia, C. Liu and S. Luo, *Appl. Catal., B*, 2018, **239**, 545–554.
- 244 X. Xie, N. Zhang, Z.-R. Tang, M. Anpo and Y.-J. Xu, *Appl. Catal., B*, 2018, **237**, 43–49.

