Recent Progress in Metal-Organic Frameworks for Applications in Electrocatalytic and Photocatalytic Water Splitting

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1. Introduction

The global demand for energy has increased rapidly and continuously in recent decades due to the quickly expanding human population and industrialization; as a result, there has been a significant increase in the utilization of traditional fossil fuels, which has caused severe environmental problems, such as the greenhouse effect, air pollution and water pollution. The quickly expanded energy consumption has resulted in major concerns about energy crises due to the limited fossil fuels resources. For a sustainable future, the development of alternative energy material that is clean and sustainable is highly desirable but remains a major challenge. Among the various energy carriers (materials), hydrogen is one of the most ideal and cleanest energy materials for the future due to its high gravimetric energy density (120 vs. 44 MJ kg⁻¹ for gasoline), high combustion efficiency, non-toxicity, clean exhaust products, and renewable and storable nature. During the past two decades, tremendous attention has been given to the field of hydrogen energy by researchers and governments around the world. However, the success of the hydrogen economy is strongly determined by the availability of useful routes for the large-scale generation of hydrogen. Currently, the production of hydrogen mainly relies on steam reforming and partial oxidation of fossil fuels (natural gas or other hydrocarbons), causing concerns about serious CO₂ emissions and limited natural resources.¹⁻³ Water, one of the most abundant resources on earth, is composed of hydrogen and oxygen atoms. Water splitting is one of the most effective ways to produce hydrogen. Currently, the production of hydrogen mainly relies on steam reforming and partial oxidation of fossil fuels (natural gas or other hydrocarbons), causing concerns about serious CO₂ emissions and limited natural resources.¹⁻³ Water, one of the most abundant resources on earth, is composed of hydrogen and oxygen atoms. Water splitting (H₂O → H₂ + 1/2O₂) consists of two half reactions, known as the oxygen evolution reaction (OER) and the hydrogen evolution reaction (HER). However, these reactions have sluggish kinetics and require catalysts. In the electrochemical process, the OER and the HER are generally catalyzed by precious metal (Ir/Ru and Pt, respectively).
materials to achieve favorable reaction kinetics. Unfortunately, noble metals suffer from low abundance and high cost, hindering their large-scale use in water electrolysis. To ensure sustainable hydrogen production, it is of great importance to seek earth-abundant alternatives to precious metal-based catalysts with excellent activity and robust stability. For example, several electrocatalysts composed of earth-abundant elements (e.g., Fe, Co and Ni) were found to be promising alternatives to precious catalysts, achieving high OER and HER activity. In addition, some carbon-based or heteroatom-doped carbon materials have been evaluated as innovative options as electrocatalysts for the OER and HER. The availability of different carbons (nanotubes, graphene, etc.) with adjustable compositions has markedly increased the number of candidates for OER/HER electrocatalysis. Solar-driven H₂ generation from water using semiconductor-based photocatalysts is another attractive route to solve the energy and environmental problems. To date, a number of metal oxide-based photocatalysts have been demonstrated to be effective for water splitting under UV light irradiation. In particular, TiO₂ has been reported as a benchmark for the UV-light-driven water splitting reactions due to its good photo-stability, low toxicity, large abundance and low cost. Unfortunately, TiO₂ has a large band gap of 3.2 eV, which can only be used in the UV light range, which includes only 5% of all solar energy (solar conversion efficiency in UV light is only 2% compared with 16% when visible light up to 600 nm can be utilized). Thus, the development of new photocatalysts with high photocatalytic activity under visible light irradiation is one of the most attractive research topics in photocatalytic water splitting. In addition, in the material composition, the activity of catalysts for electrocatalytic/photocatalytic water splitting relies heavily on the morphology of the catalyst. Thus, optimizing the catalyst composition and morphological structure is of critical importance to achieve highly efficient hydrogen production from water splitting.

Metal-organic frameworks (MOFs) are a new class of porous materials with unique electronic, optical and catalytic properties. In addition, they can be used as precursors for the fabrication of various metal, metal oxide-carbon composites or pure carbon materials with rich morphological structures and versatile properties. In applications as electrocatalysts or photocatalysts or their precursors, MOFs offer several advantages, such as high design flexibility, tunable pore channels, large surface-to-volume ratios, flexibility to be functionalized with various ligands and metal centers, and rich compositions. The metal centers separated by organic linkers in MOFs can be considered as quantum dots; consequently, short diffusion lengths of the charge carriers can be achieved during the electrocatalytic and photocatalytic reactions. The specific surface areas and band gaps of MOFs can be tailored by tuning the organic ligands and/or metal centers, so their electrocatalytic and photocatalytic activities can be tailored to maximize their performance. In recent years, MOFs have been exploited directly as photocatalysts or as their precursors for hydrogen generation from water splitting, the degradation of organic pollutants and the reduction of CO₂ into useful fuels. Recently, MOF-based materials have also proved to be particularly suitable for electrocatalytic water splitting. In the last five years, tremendous efforts have been made to apply MOFs as photocatalysts and electrocatalysts for water splitting, and interest in this research field is projected to continue increasing. Thus, a review of the recent advances and challenges of MOF-based materials in photocatalytic and electrocatalytic water splitting is highly desirable.

Herein, the recent development of MOF-based materials for electrocatalytic and photocatalytic water splitting reactions is presented. Several critical factors that determine the activity for water splitting reactions are summarized, and strategies related to the design of catalysts are emphasized. Major challenges in the fields of photocatalytic and electrocatalytic water splitting are highlighted, and some perspectives from the current progress in the development of MOF-based catalysts are given. Directions of the future research are also presented, with emphasis on achieving the desired MOF functionality and establishing structure-property relationships to identify and rationalize the factors that determine the catalytic performance. This paper aims to provide a comprehensive review of the recent progress in this dynamic field, as well as some guidelines for the further development of highly efficient photocatalysts and electrocatalysts based on MOFs for water splitting.
2. Fundamentals of Water Splitting Reactions

2.1. Electrocatalytic Water Splitting

2.1.1. Basic Principles

Electrocatalytic water splitting involves two half reactions (OER and HER), and the mechanistic schemes of the OER and HER have been proposed in the literature. The OER, which is a four-electron process, is more complex than the HER and involves several surface-adsorbed intermediates. In the following section, we mainly focus on the mechanistic study of the OER while that of the HER is described only briefly.

In the HER, the chemical adsorption and desorption of H atoms are competitive processes. A good HER catalyst should have a bond with the adsorbed H* (the asterisk indicates a bond to the catalyst surface) that is sufficiently strong to enable the proton-electron-transfer process and also sufficiently weak to ensure easy bond breaking and release of the produced H2 gas. The change in the Gibbs free energy for H* adsorption on an electrocatalyst surface (ΔG*) can be applied to evaluate both H* adsorption and H2 desorption using the HER free energy diagram. The optimal ΔG*H should be zero, under which condition the HER reaches the maximum rate. More importantly, a “volcano curve” correlation has been proposed between the experimental HER activity (HER exchange current density) and the quantum chemistry-derived ΔG*H for various catalyst surfaces. As a result, the relationship between the nature of the electrocatalyst surface and the HER kinetics can be established.

The OER pathways, in acidic or alkaline media, include elementary steps that differ according to different mechanisms, yet all involve the adsorption/desorption of intermediates, such as HO*, O* and HOO*.[55–57] The free adsorption energies of the OER intermediates at selected potentials on Pt (111) and Au (111) and some other metals were studied in acidic environment by Rossmeisl et al. using density functional theory (DFT) calculations.[58] The most difficult step in the OER is the formation of HOO* on the metal surface by splitting water on an adsorbed oxygen atom (O*). This step is downhill in free energy at high electrode potentials. At lower potentials, although water can dissociate to O*, the OER is initiated only on the oxidized surface, which makes this step slower than the O* formation process. In other words, the formation of OOH* from O* is uphill for the OER at the equilibrium potential of 1.23 V vs. reversible hydrogen electrode (RHE). Applying a voltage to move the potential positively away from 1.23 V (the difference defined as the overpotential) is thus necessary for spontaneous OER. The calculations show that the OER on Pt and Au surfaces should start at approximately 1.8 V. Simple linear relations between the stability of different intermediates and OER activity were found when the analysis was extended to other metals, which suggests that the oxygen adsorption energy is a good descriptor of the capability of a metal-based electrocatalyst for the OER.[58]

In addition to metallic catalysts, the OER mechanism on oxide catalysts has also been studied using computational methods. Rossmeisl and co-workers investigated the trends between the stability of different intermediates and OER activity. For oxide-based electrocatalysts, the intrinsic catalytic activity often related to the material composition, mixed valence states of the compositional cations (redox couples), crystal structure, metal-oxygen bond energy, oxygen vacancy concentration, electronic conductivity and charge transfer capability.[56,62–65] The number of active sites can be increased by building high-surface-area structures, tuning the morphology and creating nanostructured catalytic systems. Compositing with other catalytic materials or conductive supports can result in hybrids with enhanced activity and more active sites, which is sometimes known as the synergistic effect. The most effective methods to maximize the HER/OER activity include tailoring the surface and/or bulk properties (by the selection of cations and anions), optimizing the morphology (by the use of advanced synthetic procedures), enhancing the charge transfer process (by the functional modification of the surface electronic structure) and forming composite or hybrid catalysts. These methods may produce more active sites for HER/OER and ideal pathways for the transportation of reactants and gaseous products (i.e., hydrogen and oxygen). The strategies for enhancing electrocatalytic activity are not limited to oxides and can be, in principle, applied to other types of electrocatalysts. Additionally, researchers often take advantage of several combined strategies to improve the efficiency of electrocatalysts in the HER/OER.

The morphology and microstructure are crucial characteristics for electrocatalysts because they have a direct correlation with the number of active sites and, therefore, the catalytic activity.[66,67] For example, a simple self-template
strategy was developed to fabricate hollow Co-based bimetallic sulfide (M\(_4\)Co\(_3\)S\(_4\), M = Zn, Ni and Cu) polyhedra from homogenous bimetallic MOFs.[66] The combination of polyhedral morphology, hollow structure, homo-incorporation of a second metal element and high Brunauer-Emmett-Teller (BET) surface area significantly enhanced the HER activity of Co\(_3\)S\(_4\). The hollow Zn\(_{16}\)Co\(_{70}\)S\(_4\) exhibited the highest catalytic activity among the four electrocatalysts, indicating that cation selection is very important to achieve high electrocatalytic activity for water splitting.

In addition, nanostructured electrocatalysts generally benefit from increased specific surface area and, therefore, have more active sites for the electrocatalysis, which can be tailored by the preparation methods and annealing conditions.[68,69] For example, Shi et al. utilized an in situ carburization method to prepare MoC encapsulated by a graphitized carbon shell (nanoMoC@GS) electrocatalyst from a Mo-based MOF.[68] The nanoMoC@GS showed favorable activity in acidic media as an electrocatalyst for HER, which stemmed from the synergistic effects of the ultrafine MoC, ultrathin and conductive GS, high porosity and high surface area.[68]

Other methods to improve the surface area, such as synthesizing nanoparticles (NPs) and combining NPs with high-surface-area supports, have been used to enhance the activity of electrocatalysts for water splitting.[70,71] For example, Li et al. synthesized a nitrogen-doped Fe/Fe\(_3\)C@graphitic layer/carbon nanotube hybrid (Fe/Fe/C@NGL-NCNT) using MIL-101 (Fe) MOF as the precursor.[70] This Fe/Fe/C@NGL-NCNT hybrid showed superior OER activity and stability compared with the commercial Pt/C, which may originate from the abundant active sites and the synergistic effect of the unique architecture.

The charge transfer capability is also essential for achieving high electrocatalytic activity for water splitting, and the coupling of some functional materials, such as reduced graphene oxide (RGO), to MOF-based electrocatalysts can improve the charge transfer capability (conductivity).[72,73] For example, Tang et al. used a simple pyrolyzing method to synthesize a porous Mo-based hybrid from a polyoxometalate-based MOF and graphene oxide (POMOFs/GO), which showed improved performance for the HER.[73]

### 2.2. Photocatalytic Water Splitting

#### 2.2.1. Mechanism and Reaction Steps

Studies on splitting water into hydrogen and oxygen using light (photons) originated from the discovery of the Honda-Fujishima effect in 1967. Water splitting using photocatalysts has since been widely investigated.[74–78] Previous reviews of water splitting using semiconductors as photocatalysts have demonstrated the basic principles of the water splitting process.[76–78] The electrons in the valence band (VB) of the photocatalyst are transferred to the conduction band (CB), and holes are left in the VB after absorbing UV and/or visible light, creating electron-hole pairs. The photogenerated electron-hole pairs can induce redox reactions similar to water electrolysis. Specifically, water molecules are reduced by the electrons to generate H\(_2\) and are oxidized by the holes to produce O\(_2\), completing the water splitting reactions.

Water splitting into H\(_2\) and O\(_2\) is an energetically uphill reaction with a standard Gibbs free energy change (\(\Delta G\)) of +237 kJ mol\(^{-1}\) (corresponding to 1.23 eV). Therefore, the band gap of the photocatalysts and the edges of the CB and VB must be suitable for water splitting. The top level of the CB should be more negative than the redox potential of H\(^+\)/H\(_2\) (0.0 V vs. normal hydrogen electrode (NHE)), while the top level of the VB should be more positive than the redox potential of O\(_2\)/H\(_2\)O (1.23 V vs. NHE).[74] The theoretical minimum band gap for water splitting is therefore 1.23 eV, which is equivalent to a light wavelength of approximately 1100 nm. However, not all semiconductors meet the requirements for water splitting. For metal oxide-based photocatalysts, the VB mainly consists of O 2p orbitals, and the top level of the VB is much higher than 1.23 V vs. NHE. Therefore, the oxidation-reduction potentials (ORPs) of O\(_2\)/H\(_2\)O and H\(^+\)/H\(_2\) are positioned between the top level of the VB and the bottom level of the CB. Higher photon energy than the band gap of the photocatalyst is needed due to an activation barrier in the charge transfer process in the water splitting reactions. At the same time, the much wider band gaps of these materials make them only photoactive in UV light. As approximately 50% of the solar spectrum consists of visible photons (400 < \(\lambda\) < 800 nm), it is critical to develop active photocatalysts with high activity under visible light for photocatalytic water splitting.

The development of photocatalysts is very important to enable water splitting with visible light. The main steps in the photocatalytic water splitting reactions should be tailored to meet the requirements for photocatalysts capable of water splitting. There are three steps in the photocatalytic water splitting reactions, which has been demonstrated in some informative reviews.[76–78] The first step is the formation of electron-hole pairs by incident photons. When the energy of incident light is greater than the band gap energy, the electrons in the VBs can be excited and transferred into the CB. Meanwhile, holes are generated in the VB. However, the band structure is only a thermodynamic requirement. Other factors, such as charge separation, mobility and the lifetime of photogenerated electrons and holes can also affect the photocatalytic activity for water splitting. The second step is charge separation and diffusion to the catalyst surface without recombination of the photogenerated carriers, which is drastically affected by the crystal structure, particle size and crystallinity of the photocatalyst.[79,80] A higher crystallinity can lead to superior charge migration efficiency because the defects in a photocatalyst with lower crystallinity act as recombination centers for the photogenerated electron-hole pairs, which decreases the photocatalytic activity. A smaller particle size of the photocatalyst also suppresses the possibility of electron-hole pair recombination. The final step is the reduction and oxidation of surface-adsorbed species by the photogenerated electrons and holes to generate H\(_2\) and O\(_2\), respectively. In this step, the surface active sites of the photocatalyst play vital roles in efficient water splitting. Co-catalysts, such as Pt, are usually loaded onto the photocatalyst surface as active sites to reduce the activation energy for the HER.[83] These processes affect the overall efficiency of water splitting based on a semiconductor-based photocatalyst.
2.2.2. Factors to Determine the Photocatalytic Activity

The main factors that determine the photocatalytic activity of the photocatalysts for water splitting include the band gap energy/visible light absorption capability, active sites/co-catalysts and charge transfer/separation efficiency. Covalent modification is a method to reduce the band gap energy of photocatalysts. For example, the band gap of MOFs can be reduced by diazo coupling with amino-substituted ligands and other molecules.[82] The photocatalytic activity of these modified MOFs corresponds to a red shift of the absorption edge, suggesting that the band gap energy/visible light absorption capability plays a vital role in the photocatalytic activity.

The incorporation of active co-catalysts is an effective way to increase the number of active sites for water splitting.[83–87] For example, the well-defined cages of MIL-101(Cr) MOF were used to engage the molecules of a high-valent di-n-oxo dimanganese catalyst with high activity for photo-electrochemical (PEC) water oxidation and the incorporation of MnTD ([terpy] Mn(μ-O)2Mn(terpy)]3+: terpy: 2,2′,6′,2″-terpyridine) improved the turnover number of MIL-101(Cr) more than 20-fold while maintaining the initial high rate in the PEC water oxidation reaction.[81] In another study, Hansen and Das found that MnTDC=MIL-101(Cr) showed superior activity to MIL-101(Cr) and MnO2 catalysts for the OER.[84] These studies suggested that the incorporation of active co-catalysts can greatly enhance the photocatalytic activity for water splitting, and the selection and incorporation method of the co-catalysts should be optimized.

The charge transfer/separation efficiency of the photocatalyst plays a critical role in photocatalytic water splitting. The construction of heterojunctions is an effective way to enhance the charge transfer/separation capability of electron-hole pairs.[88,89] For example, a MOF-derived Co3O4/TiO2 composite photocatalyst with 2 wt.% Co loading and a p-n heterojunction, exhibited a much higher hydrogen evolution rate than the conventional Co3O4/TiO2 nanocomposite (= 7-fold enhancement).[88] The photocatalytic activity of MOF-based photocatalysts is determined by several crucial factors, such as the band gap energy, active sites/co-catalyst and charge transfer capability. These factors are often closely related. For example, an azocarboxylic acid can be used as an organic linker to construct a Gd-based MOF with a reduced band gap.[89] Gd-MOF has high photocatalytic activity for the HER due to its high visible light absorption capability. The addition of Ag co-catalyst improved the HER activity of Gd-MOF by providing more active sites and improving the charge transfer capability.

3. Recent Advances in MOF-Based Catalysts for Water Splitting

Because of the many outstanding features of MOFs, such as tunable pore channels, high specific surface area, easy tailoring of the material composition, rich morphological structure, and capability to act as precursors for the preparation of various metal/metal oxide/carbon composites and carbon materials of various properties, during the past five years, the applications of MOFs as catalysts or the precursors of catalysts for electrocatalytic and photocatalytic water splitting reactions for hydrogen generation have been extensively exploited. Both the direct application of MOFs for water splitting and application as a precursor for metal/metal oxide/carbon composites or porous carbon materials (by leaching of the metal/metal oxide from the composites), which were then applied as electrocatalysts or photocatalysts, have been reported. Additionally, MOFs were studied as catalysts for both the OER and HER, and the reactions were conducted in acidic and alkaline electrolytes.

3.1. Electrocatalytic Water Splitting

For photocatalytic water splitting, the direct application of MOFs as electrocatalysts was first reported in 2011 by Nohra et al., who pioneered the use of polyoxometalate-based MOFs (POMOFs) for the HER.[90] The structural properties were investigated but their electrocatalytic activity was only briefly studied and the efficiency of POMOFs to replace Pt catalyst for the HER was not clearly demonstrated. In 2015, Qin et al. reported a type of POMOFs called [TBA][r-PMo12V4O41(OH)3(Zn4)[BTB]1/3: xGuest (NNU-500, BTB = benzene tribenzoate, TBA+ = tetrabutylammonium ion) as an ultrastable electrocatalyst for the HER.[91] It displayed a Tafel slope of 96 mV dec−1 and an overpotential of 237 mV at a current density of 10 mA cm−2 (a metric associated with solar fuel synthesis), which was inferior to Pt/C (Tafel slope of 30 mV dec−1 and overpotential of 52 mV at 10 mA cm−2).

Very recently, Dai et al. demonstrated MoSx nanosheets to the MOFs dramatically enhanced the HER activity due to the improved electron transport, the increased number of active sites and the favorable delivery of local protons in the Zr-MOF structure. By optimizing the MoSx amount, the MoSx-MOF composite with a Mo/Zr ratio of 0.5 displayed remarkable HER activity, with a Tafel slope of 59 mV dec−1, which was only slightly higher than that of Pt/C (32 mV dec−1).[91]

In the study of MOFs as precursors for the preparation of electrocatalysts for water splitting reactions, Chaikittisilp and co-workers were the first to use a Co-based MOF (zeolitic imidazolate framework-9, ZIF-9) as a precursor for the preparation of a nanoporous CoOx-C hybrid as an electrocatalyst for the OER.[94] The conversion of ZIF-9 to the CoOx-C hybrid is shown in Figure 1a. As depicted in Figure 1b, for the OER, the Z9-700-250 and Z9-800-250 electrocatalysts exhibited more negative onset potentials and higher current densities than Z9-900-250 and Pt/carbon black. These results indicated that the Z9-800-250 hybrid is a promising electrocatalyst for the OER. Very recently, Aijaz et al. reported a highly active electrocatalyst for the OER comprising core-shell Co@Co3O4 NPs embedded in CNT-grafted N-doped carbon-polyhedra, which was obtained by the pyrolysis of a Co-based MOF in H2 atmosphere and a subsequent controlled oxidative calcination.[95] This electrocatalyst displayed an overpotential of 410 mV at 10 mA cm−2, comparable to RuO2, which has been demonstrated as the benchmark electrocatalyst for the OER.

Although nanoporous carbon was successfully synthesized from MOFs in 2008[96] and was widely used in the oxygen reduction reaction (ORR).[97–99] the direct use of nanoporous carbon derived from MOFs in water splitting was demonstrated only...
Among the various parameters, the morphology of the electrocatalysts can be tuned through the preparation method. For example, a Zn-based MOF, MOF-5, was prepared by an ionic liquid (IL)-based method and displayed a distinct flower-shaped morphology with a diameter of approximately 10 µm, very different from the regular cubic structure of MOF-5 prepared by traditional methods. The as-prepared MOF-5(II) displayed superior activity for the HER compared to cubic MOF-5 due to the enhanced oxygen adsorption reaction of the hydrogen atoms.\(^\text{[101]}\)

In addition to the direct use of MOFs in electrocatalytic water splitting, MOFs have also been applied as precursors for the synthesis of electrocatalysts with controlled particle sizes and morphologies.\(^\text{[102–105]}\) The morphology of the electrocatalysts derived from MOFs precursors can be tailored through the choice of the type of MOF and the temperature and atmosphere used for the subsequent calcination. For example, a CoP electrode with a concave polyhedrons (CPHs) morphology was synthesized by topological conversion using ZIF-67 polyhedrons as the precursor.\(^\text{[102]}\) The morphology of the CoP CPHs is shown in Figure 2a. CoP NPs synthesized from direct calcination of Co(NO\(_3\))\(_2\) show a different morphology (Figure 2b). For the HER, the CoP CPHs electrocatalyst showed a low overpotential of 133 mV at 10 mA cm\(^{-2}\), while the CoP NPs had an overpotential of 187 mV under the same conditions. In addition, the Tafel slopes of the CoP CPHs and CoP NPs are 51 and 63 mV dec\(^{-1}\), respectively. The electrocatalytic performance of the porous CoP CPHs is superior to most of the reported non-noble-metal-based catalysts, such as CoP microspheres and FeP nanosheets, for the HER.\(^\text{[106–107]}\)

In addition to CoP CPHs, electrocatalysts with other controlled morphologies, such as nano-octahedrons, spindle-like 3D structures and porous nanocages, were also synthesized from MOFs and were investigated as electrocatalysts for the OER and HER, showing attractive activity.\(^\text{[108–110]}\) For example, Wu et al. introduced a MOF-assisted strategy for the synthesis of MoC\(_x\) nano-octahedrons as electrocatalysts for the HER.\(^\text{[108]}\) The MoC\(_x\) nano-octahedrons (Figure 3a) exhibited superior electrocatalytic activity and stability compared to irregular-shaped MoC\(_x\) NPs with a similar composition (Figure 3b) in both acidic and basic solutions. The overpotential of MoC\(_x\) nano-octahedrons was 87 and 92 mV at 1 mA cm\(^{-2}\) in acidic and basic solutions, respectively, while it was approximately 230 mV for the irregular-shaped MoC\(_x\) NPs in acidic and basic solutions.

In another study, MOF-derived Ni-Co-based metal oxides with nanocage morphology were compared with a catalyst with a nanocube morphology and the same composition and comparable surface area (~31 m\(^2\) g\(^{-1}\)) for water splitting.\(^\text{[110]}\) Ni-Co Prussian-blue-analog (PBA) nanocubes were derived from the Ni-Co-PBA nanocubes as precursors and were converted to Ni-Co mixed oxides by calcination in air. As shown in Figure 4a,b, the Ni-Co mixed oxides inherited the nanocage morphology of the Ni-Co PBA particles but had rougher surfaces than the pristine Ni-Co mixed oxide nanocubes. As shown in Figure 4c,d, the Ni-Co mixed oxide nanocubes exhibited lower onset potential, lower Tafel slope and higher current density than the nanocubes for the OER. The enhanced OER activity of the Ni-Co mixed oxide nanocubes was assigned to the hollow and porous nanocage morphology.
3.1.2. Constructing Hybrids/Composites

Because of the large variation of organic moieties in MOFs, highly nanoporous carbon materials may be formed by thermal carbonization of MOFs in inert atmospheres. Such MOF-derived nanoporous carbon materials usually exhibit exceptionally high surface areas, which can form strong coupling with metal oxides/metal to enhance the electrocatalytic activity and stability for water splitting. Thus, many investigations have attempted the synthesis of carbon-metal/metal oxide composites from MOFs as electrocatalysts for the HER and OER. For example, hybrid porous nanowire arrays (NAs) composed of strongly interacting Co₃O₄ and carbon (Co₃O₄C-NA) with a high specific surface area of 251 m² g⁻¹ and a large carbon content of 52.1 wt.% were successfully prepared by a facile carbonization of MOF grown on Cu foil, and displayed superior performance as the working electrode for OER without additional substrates or binders. A low onset potential of 1.47 V vs. RHE and a stable current density of 10 mA cm⁻² at 1.52 V vs. RHE in 0.1 M KOH for 30 h were achieved. Co₃O₄C-NA also showed a much lower Tafel slope of 61 mV dec⁻¹ than IrO₂/C (87 mV dec⁻¹) in 1 M KOH. For comparison, the carbon-free counterpart (Co₃O₄-NA), prepared by the calcination of Co₃O₄C-NA in air, which exhibited a similar porous NA structure and a cubic spinel phase, displayed a higher onset potential of 1.50 V vs. RHE, a larger operating potential of 1.64 V vs. RHE at 10 mA cm⁻² and a much higher Tafel slope of 123 mV dec⁻¹. Co₃O₄C-NA delivered superior OER activity and stability compared with the state-of-the-art noble-metal electrocatalyst.

The strong coupling was utilized for the development of a MoS₂-based composite (MoS₂/3D-NPC) as the electrocatalyst for the HER, in which the MoS₂ nanosheets grew in situ in the nanopores of 3D nanoporous carbon (3D-NPC) derived from MOF and showed much better performance than the respective components (MoS₂ NPs, 3D-NPC) of the catalyst, as well as the physically mixed MoS₂ NPs and NPC composite (MoS₂+C-3D-NPC).

The MoS₂/3D-NPC composite displayed a small overpotential of 210 mV at 10 mA cm⁻² for the HER, better than those of 3D-NPC (≈300 mV), MoS₂ NPs (≈250 mV) and MoS₂ + 3D-NPC (≈250 mV) as shown in Figure 5a. The Tafel slope of the MoS₂/3D-NPC composite reached 51 mV dec⁻¹, in contrast to 95 mV dec⁻¹ for the MoS₂ NPs as shown in Figure 5b. The high electrocatalytic activity of the MoS₂/3D-NPC electrocatalyst was attributed to the efficient charge transfer in the HER due to the more exposed active sites and robust interaction between the MoS₂ and the MOF-derived conductive carbon when the MoS₂ nanosheets were grown in the pores of the 3D-NPC.

In another study, Lu et al. reported the development of a core (Au NP, ≈50–100 nm)-shell (Zn-Fe-C, ≈30–60 nm) composite prepared by direct pyrolysis of a Zn-Fe-MOF shell coated on an Au NP in an inert atmosphere. The core-shell catalyst (Au@Zn-Fe-C) displayed a low onset potential of 0.08 V vs. RHE in 0.5 M H₂SO₄, which was much more positive than those of Au (≈0.225 V) and Zn-Fe-C (≈0.292 V) but slightly more negative than that of commercial Pt/C (≈0.006 V). The Tafel slope of Au@Zn-Fe-C was 130 mV dec⁻¹, which was lower than the slopes of Au (167 mV dec⁻¹) and Zn-Fe-C (271 mV dec⁻¹). The onset overpotential and lower Tafel slope contributed to the more favorable HER kinetics of Au@Zn-Fe-C, demonstrating a synergistic effect between the Au NP core and Zn-Fe-C shell.

3.1.3. Functional Modification

The charge transfer capability of the electrocatalyst plays a critical role in determining the activity for OER and HER. For MOF-based catalysts, the addition of functional ions and/or ligands can enhance the charge transfer capability. For example, Wang et al. developed a new electrocatalyst by assembling Co ions and benzimidazolate ligands into a MOF (Co-ZIF-9) for the OER with high proton transfer capability. Co-ZIF-9 was effective for the electrocatalytic OER, and its turnover frequency (TOF) reached ≈1.76 × 10⁻³ s⁻¹, which
is similar to the active Co-based electrocatalyst reported in the literature.\cite{121} Wang et al.’s work made an important step in water splitting chemistry by integrating the redox-active metal centers and organic motifs into a MOF structure.

In addition to the design of MOFs, some carbon-based functional additives were incorporated into MOF-based catalysts to enhance the charge transfer capability.\cite{122,123} For example, GO was used to modify the MOFs to enhance the charge transfer capability and improve the electrocatalytic activity for water splitting.\cite{123} The GO-incorporated Cu-MOF composite displayed good performance as a bifunctional catalyst for the HER and OER in 0.5 M H₂SO₄.\cite{123} The (GO 8 wt.%) Cu-MOF exhibited the highest activity for the HER. The onset overpotential for the HER was 202 mV for Cu-MOF and decreased from 123 to 87 mV as the GO amount increased from 2 to 8 wt.%. For the OER, the Tafel slope was 65 mV dec⁻¹ for (GO 8 wt.%) Cu-MOF and 89, 81 and 61 mV dec⁻¹ for Cu-MOF, (GO 2 wt.%) Cu-MOF and Pt/C, respectively. The improved activity of the GO-MOF composite for water splitting in acidic solution was attributed to the enhanced charge transfer capability and synergistic effects of GO and MOF.

Functional carbon-based materials were also added to MOF-based materials in the precursor stage for the preparation of

Figure 4. TEM images of the as-prepared Ni-Co oxide nanocages (a) and porous nanocubes (b); Polarization curves (c) and Tafel plots (d) of the Ni-Co mixed oxide nanocages and porous nanocubes for the OER. Reproduced with permission.\cite{110}

Figure 5. a) Polarization curves and b) Tafel plots of the MoS₂/3D-NPC composite, 3D-NPC, MoS₂ NPs, mixed MoS₂+3D-NPC and Pt. Reproduced with permission.\cite{118} Copyright 2015, Royal Society of Chemistry.
highly active MOF-derived electrocatalysts. For example, Tang et al. synthesized an active electrocatalyst derived from a POMOFs/GO composite for the HER. The introduction of GO to POMOFs improved the conductivity and the HER activity of MoO$_2$@PC-RGO and functioned as a support for the formation of a closely connected network. This hybrid presented superior activity for the HER in acidic media due to the synergistic effects among the MoO$_2$ NPs, PC and RGO substrates. MoO$_2$@PC-RGO initiated H$_2$ evolution near its thermodynamic overpotential (0 mV), which was similar to that of the Pt/C catalyst, while MoO$_2$@PC had an onset overpotential of 66 mV. The overpotentials at 10 mA cm$^{-2}$ were 38 mV for Pt/C, 64 mV for MoO$_2$@PC-RGO and 136 mV for MoO$_2$@PC. Tafel slopes of 30, 41 and 60 mV dec$^{-1}$ were obtained for Pt/C, MoO$_2$@PC-RGO and MoO$_2$@PC in 0.5 M H$_2$SO$_4$.

In another work, rGO was incorporated with CoP to enhance the charge transfer capability of CoP for the HER and OER in alkaline media. This layered composite was prepared by pyrolysis and phosphating processes with rationally designed sandwich-type ZIF-67 MOF/GO as a template and precursor, as shown in Figure 7. The MOF-derived porous CoP nanostructure guaranteed a large quantity of exposed active sites, and the close contact between CoP and rGO contributed to a continuous conductive network, which was beneficial for the electron transfer process. For the HER, CoP/rGO-400 was more active than rGO and CoP due to the synergistic effect between the CoP and rGO in the composite. In 1 M KOH, a low Tafel slope of 38 mV dec$^{-1}$ was achieved with the CoP/rGO-400 electrocatalyst, comparable to that of Pt/C (36 mV dec$^{-1}$) and much lower than CoP (60 mV dec$^{-1}$). For the OER in the same alkaline solutions, CoP/rGO-400 displayed a Tafel slope of 66 mV dec$^{-1}$, which was superior to CoP, rGO and even the state-of-the-art IrO$_2$. These contributions shed light on the rational design of a series of RGO-incorporated MOF-based electrocatalysts for the OER and HER.

3.1.4. Intrinsically Conductive MOFs

MOF is typically not conductive and may not be a great choice for electrocatalysis-based applications. As such, one of the most challenging and rewarding endeavors in this field is to synthesize porous MOFs with good charge mobility and conductivity. Recently, the development of new type MOFs with enhanced intrinsic conductivity has attracted increasing attention. In 2009, Takaishi et al. reported one of the first conductive MOFs, Cu[$\text{Cu}$(pdt)$_2$] (pdt$^-$=2,3-pyrazinedithiolate), and its electrical conductivity was $6 \times 10^{-4}$ S cm$^{-1}$ at 300 K. However, this kind of MOF collapsed upon desolvation. Thus, it is critical to develop alternative MOFs with high conductivity and structural stability for the potential electrocatalysis application.

Dinca and co-workers have made great contributions to the development of conductive MOFs for the potential use in electrocatalysis-based applications. For example, Sun et al. treated Mn$_2$$^{2+}$ with 2,5-disulfhydrylbenzene-1,4-dicarboxylic acid (H$_4$DSBDC) and obtained isolated Mn$_2$(DSBDC). This MOF was a thiolated analogue of Mn$_2$(DOBDC) MOF derived from 2,5-dihydroxybenzene-1,4-dicarboxylic acid (H$_4$DOBDC). The porous Mn$_2$(DSBDC) with one-dimensional (1D) (–Mn–S–)$_\infty$ chains showed a high surface area (978 m$^2$ g$^{-1}$) and high charge mobility (0.01 cm$^2$ V$^{-1}$ s$^{-1}$) similar to those of the most common organic semiconductors. Although Mn$_2$(DSBDC) displayed a relatively high charge mobility, its conductivity was rather low (3.9 $\times 10^{-13}$ S cm$^{-1}$ at 297 K). The replacement of d$^5$ Mn$^{2+}$ by d$^6$ Fe$^{3+}$ centers introduces high energy, loosely bound minority-spin carriers and then enhances the conductivity. They found that the bulk electrical conductivity values of both...
Fe$_2$(DSBDC) and Fe$_2$(DOBDC) were ≈6 orders of magnitude higher than those of the Mn$^{2+}$ analogues, Mn$_2$(DOBDC) and Mn$_2$(DSBDC), which was attributed to the loosely bound Fe$^{2+}$β-spin electron. This study can provide important insights for the rational design of conductive MOFs, highlighting in particular the advantages of iron for synthesizing conductive MOF-based materials.

Besides Fe, Cd was also demonstrated to be a promising candidate to construct conductive MOFs. Park et al. found that isostructural MOFs M$_2$(TTFTB) (M = Mn, Co, Zn and Cd; H$_4$TTFTB = tetrathiafulvalene tetrabenzoate) exhibited a striking correlation between their single-crystal conductivities and the shortest S···S interaction defined by neighboring tetrathiafulvalene (TTF) cores, which inversely correlated with the ionic radius of the metal ions.$^{[132]}$ The Cd analogue, with the largest cation and shortest S···S contact, showed the highest electrical conductivity ($2.86 \times 10^{-4}$ S cm$^{-1}$) that was 72 times higher than that of Zn$_2$(TTFTB) ($3.95 \times 10^{-6}$ S cm$^{-1}$). Mn$_2$(TTFTB) and Co$_2$(TTFTB), which display intermediate S···S distances between those observed in the Zn and Cd analogues, also showed intermediate conductivity values of $8.64 \times 10^{-5}$ and $1.49 \times 10^{-5}$ S cm$^{-1}$, respectively, both tracking inversely with increasing S···S distance.

Highly conductive two-dimensional (2D) MOFs made from nitrogen-based ligands were reported in 2014 by Sheberla et al.$^{[133]}$ They found that the reaction of NiCl$_2$ with hexaaminotriphenylene (H$_6$HATP) in aqueous NH$_3$ solution led to the isolation of a new MOF, Ni$_3$(HITP)$_2$ (HITP = 2,3,6,7,10,11-hexaiminotriphenylene). Ni$_3$(HITP)$_2$ films grown on a quartz substrate displayed a conductivity of 40 S cm$^{-1}$ at room temperature, while pellets of the same material showed a bulk conductivity of 2 S cm$^{-1}$. The isostructural material made from Cu$_{11}$, Cu$_2$(HITP)$_2$, also displayed a high bulk conductivity of 0.2 S cm$^{-1}$ at room temperature.

The large bulk conductivity of HITP-based materials enabled the application in chemiresistive sensing for ammonia vapor.$^{[134,135]}$ Ni$_3$(HITP)$_2$ functioned as reversible chemiresistive sensors, which was capable of detecting sub-ppm levels of ammonia vapor.$^{[134]}$ It was found that the chemiresistive response can be altered by the choice of metal node and Ni$_3$(HITP)$_2$ was unresponsive to ammonia, suggesting that the copper sites are critical for ammonia sensing. These studies highlight the utility of 2D conductive MOFs in the production of tunable functional materials, such as chemical sensors.

Very recently, Miner et al. investigated an intrinsically conductive Ni$_3$(HITP)$_2$ MOF as an active, well-defined and tunable electrocatalyst toward the ORR in alkaline solutions.$^{[136]}$ Ni$_3$(HITP)$_2$ exhibited ORR activity competitive with the most active Pt-free electrocatalysts due to the combination of high crystallinity of MOFs, the physical durability and electrical conductivity of graphitic materials, and the diverse, well-controlled synthetic accessibility of molecular species. In addition, the Ni$_3$(HITP)$_2$ MOF with high electrical conductivity is directly used as the electrode material in electrochemical double layer capacitors (EDLCs) without conductive additives or other binders.$^{[137]}$ The performance of this MOF exceeded that of most carbon-based materials with capacity retention greater than 90% over 10000 cycles, suggesting conductive MOFs as a new generation of active materials for supercapacitors. These studies highlight the direct use of conductive MOFs in fuel cells and supercapacitors, which can shed light to their future application in the electrocatalytic water splitting.

### 3.1.5. A Brief Summary

The catalytic activities of some typical MOFs and MOF-derived electrocatalysts for the HER and/or OER are summarized in Table 1. Taking advantage of the porous structures of MOF precursors, the MOF-derived materials had various morphological features, such as polyhedrons, nanocubes, nanocages, nanooctahedrons, nanosheets and nanowires, and outperformed their NP counterparts of similar compositions for catalyzing the water splitting reactions. Remarkably, in the HER under acidic conditions, porous CoP with a CPHs structure prepared...
from Co-MOFs exhibited a >50 mV decrease in the overpotential required to reach 10 mA cm$^{-2}$ compared to CoP NPs.$^{[102]}$ As MOFs typically have abundant carbon and nitrogen sources, the carbonation of MOFs can lead to the formation of catalytically active metal-based material composites with carbon materials having finely tuned porosity, graphic degree and nitrogen content (if any). These hybrids have demonstrated good HER/OER performance, as shown in Table 1. Particularly, a strongly interacting Co$_3$O$_4$ and carbon hybrid grown on Cu foil directly catalyzed the OER with high efficiency, reaching 10 mA cm$^{-2}$ at an overpotential as low as 290 mV in 0.1 M KOH.$^{[113]}$ In addition to applying MOFs alone to synthesize electrocatalytic materials, GO and rGO were functionalized with MOFs to improve the charge transfer of the catalysts, as well as the catalytic performance. Despite the effectiveness of MOF-derived materials in the HER and/or OER, most can only promote the HER in acidic media (while industrial water splitting favors an alkaline solution) and few can catalyze the overall water splitting process (both HER and OER) in the same electrolyte. Therefore, more efforts should be made to design bifunctional electrocatalysts from MOFs with the combined knowledge of MOF structures and the mechanism of the water splitting process. More importantly, current MOF-derived catalysts can only sustain constant hydrogen or oxygen production within limited timescales, ranging from several hours to a few days. Robust MOF-derived materials that can operate on a longer timescale are thus needed to go a step further toward the real application of water splitting.

### 3.2. Photocatalytic Water Splitting

Some MOFs have shown favorable photocatalytic activity for water splitting, dye degradation, CO$_2$ reduction and Cr(VI) reduction.$^{[138–141]}$ For photocatalytic water splitting, the direct use of MOFs as photocatalysts for the HER was first reported in Table 1. Electrocatalytic activity of some typical MOF-based and MOF-derived electrocatalysts for the HER and OER.

| No. | Electrocatalyst | Target reaction | Medium | Overpotential at 10 mA cm$^{-2}$ (mV) | Tafel slope (mV dec$^{-1}$) | Durability (h) | Ref. |
|-----|-----------------|-----------------|--------|-------------------------------------|---------------------------|--------------|-----|
| 1   | (GO 8 wt%) Cu-MOF composite | OER | 0.5 M H$_2$SO$_4$ | 110 at 2 mA cm$^{-2}$ | 65 | / | [123] |
|     |                 | HER | 0.5 M H$_2$SO$_4$ | 209 at 10 mA cm$^{-2}$ | 84 | / |     |
| 2   | N-doped carbon nanotube frameworks (NCNTFs) | OER | 1 M KOH | 370 | 93 | 3 | [100] |
| 3   | Zn$_{10}$Co$_{20}$S$_{20}$ polyhedra | HER | 0.5 M H$_2$SO$_4$ | 80 | 47.5 | 60 | [66] |
| 4   | Co$_2$S$_4$ polyhedra | HER | 0.5 M H$_2$SO$_4$ | 133 | 51 | 12 | [102] |
| 5   | CoP CPHs | HER | 0.5 M H$_2$SO$_4$ | 142 | 53 | 11 | [108] |
| 6   | MoC$_4$ nano-octahedrons | HER | 0.5 M H$_2$SO$_4$ | 151 | 59 | 11 |     |
| 7   | Ni-Co mixed oxide cages | OER | 1 M KOH | 380 | 50 | 10 | [110] |
| 8   | Ni-Co mixed oxide cubes | OER | 1 M KOH | 430 | 59 | 10 |     |
| 9   | CoNi hydroxide ultrathin nanosheets | OER | 1 M KOH | 324 | 33 | 3 | [104] |
| 10  | Spindle-like Co/Fe metal oxides in N-doped porous carbon | OER | 0.1 M KOH | 390 | 72.9 | / | [109] |
| 11  | CoP$_x$ NPs embedded in N-doped carbon matrices | HER | 1 M KOH | 319 | 52 | 24 | [67] |
| 12  | Co$_3$O$_4$/C-NA | OER | 0.1 M KOH | 290 | 70 | 30 | [113] |
| 13  | MoS$_2$/3D-NPC | HER | 0.5 M H$_2$SO$_4$ | 210 | 51 | 3 | [118] |
| 14  | Au@Zn$_{10}$Fe$_{10}$C | HER | 0.5 M H$_2$SO$_4$ | 123 | 130 | 12 | [119] |
| 15  | Atomically isolated nickel species anchored on graphitized carbon | HER | 0.5 M H$_2$SO$_4$ | 34 | 41 | 25 | [117] |
| 16  | Co NPs embedded in porous N-rich carbon | HER | 1 M KOH | 370 | 76 | 10 | [115] |
| 17  | Co@Co$_3$O$_4$ encapsulated in CNT-grafted N-doped carbon polyhedra | OER | 0.1 M KOH | 410 | 54.3 | 45 | [95] |
| 18  | Layered CoP/rGO composite | HER | 0.5 M H$_2$SO$_4$ | 105 | 50 | 22 | [124] |
|     |                 | OER | 1 M KOH | 150 | 38 | 22 |     |
|     |                 | OER | 1 M KOH | 340 | 66 | 22 |     |
They demonstrated two types of Zn-based MOFs formed by terephthalate (UiO-66) and 2-aminoterephthalate ligands (UiO-66-NH$_2$) as stable photocatalysts for the HER. However, the H$_2$ generation rates of both photocatalysts were low (248 and 372 µmol h$^{-1}$ g$^{-1}$ for UiO-66 and UiO-66-NH$_2$, respectively). Thus, it is critical to incorporate a Pt co-catalyst and to optimize the Pt amount to enhance the activity of MOF-based photocatalysts.\cite{142} The optimum Pt amount (1.5 wt.%) achieved a high H$_2$ evolution rate of 500 µmol h$^{-1}$ g$^{-1}$ with a Pt/Ti-MOF-NH$_2$ photocatalyst.\cite{143} Recently, Shen et al. used an active MoS$_2$ to replace expensive Pt in the UiO-66-CdS system, and superior photocatalytic activity was achieved with the MoS$_2$/UiO-66-CdS photocatalyst.\cite{143} As shown in Figure 8a, the incorporation of CdS and UiO-66 enhanced the photocatalytic activity of both parts of the reaction, and the optimum HER activity was obtained at a UiO-66 to CdS weight ratio of 1:1. A high H$_2$ evolution rate of 25,770 µmol h$^{-1}$ g$^{-1}$ was obtained in the presence of 1 wt.% MoS$_2$, approximately 2-fold higher than that of 1 wt.% Pt/UiO-66-CdS. As shown in Figure 8b, the H$_2$ evolution rate of MoS$_2$/UiO66-CdS reached a maximum when the MoS$_2$ amount in the composite was 1.5 wt.% (32,500 µmol h$^{-1}$ g$^{-1}$). Shen et al.’s work demonstrated the advantage of using MoS$_2$ as a highly active co-catalyst to replace Pt in MOF-based photocatalysts to enhance the HER activity.

Lin and co-workers were the first to use an Fe-based MOF (MIL-101) coated with amorphous TiO$_2$ as a template to prepare Fe$_2$O$_3$@TiO$_2$ composite as a photocatalyst for the HER.\cite{144} This MOF-derived Fe$_2$O$_3$@TiO$_2$ composite exhibited superior HER activity to either Fe$_2$O$_3$ or TiO$_2$ alone and a mixture of the two. However, this composite had a low H$_2$ evolution rate of ~600 µmol h$^{-1}$ g$^{-1}$ after depositing Pt NPs as co-catalysts. More recently, Pham et al. reported a hollow Fe$_2$O$_3$@TiO$_2$-PtO$_x$ photocatalyst for the HER that was synthesized from MIL-88B as a template.\cite{145} The hollow Fe$_2$O$_3$@TiO$_2$-PtO$_x$ nanocomposite displayed an enhanced hydrogen evolution rate of 1100 µmol h$^{-1}$ g$^{-1}$ by separating the two co-catalysts (Fe$_2$O$_3$ and PtO$_x$) on two surface sides of TiO$_2$. In 2015, Bala et al. reported a MOF-derived Co$_6$O$_7$@TiO$_2$ photocatalyst with rich p-n heterojunctions for the HER.\cite{88} The Co$_6$O$_7$@TiO$_2$ composite with the optimum Co loading amount of 2 wt.% exhibited the maximum photocatalytic activity, with a hydrogen evolution rate of 7000 µmol h$^{-1}$ g$^{-1}$, which was almost 7 times greater than that of the conventional Co$_3$O$_4$/TiO$_2$ nanocomposite.\cite{88}

As demonstrated previously, the catalytic activity of photocatalysts for the HER and OER is strongly related to the band gap energy, the visible light absorption capability, the composition and quantity of active sites/co-catalysts and the charge transfer capability. In the following section, we summarize several important examples of MOFs as photocatalysts for water splitting under solar irradiation, including band gap engineering to improve the visible light response, active site/co-catalyst selection, incorporation and optimization to enhance the activity and structural evolution, functional coupling and the creation of heterojunctions to enhance the charge transfer capability.

3.2.1. Engineering the Band Gap and/or Enhancing the Visible Light Absorption Capability

Tremendous effort has been made to develop photocatalysts with high activity under visible light irradiation, and it is primarily important to tailor the band structure.\cite{34-37} A hybrid organic/inorganic compound with a MOF structure, Cu$_3$PO$_4$(C$_2$N$_3$H$_2$)$_2$OH, which was synthesized by hydrothermal route in the presence of 1,2,4-triazole as a structure directing agent, was investigated as a photocatalyst for the OER under visible light.\cite{146} This Cu-based MOF had a band gap of 2.58 eV, which ensured visible-light-driven photocatalytic activity. When Cu$_3$PO$_4$(C$_2$N$_3$H$_2$)$_2$OH was used as the photocatalyst, the O$_2$ evolution rate of ~1875 µmol h$^{-1}$ g$^{-1}$. In principle, MOFs with open metal coordination sites might be able to activate small molecules, such as water, triggering chemical reactions under mild conditions. Therefore, the water oxidation reaction based on this type of porous material should be explored.

In an attempt to enhance the activity of MOF-based photocatalysts for the HER under visible light irradiation, Toyao et al. reported a Ti-MOF-NH$_2$ photocatalyst using 2-amino-benzenedicarboxylic acid (H$_2$BDC-NH$_2$) as an organic linker (Figure 9) for the photocatalytic HER under visible light irradiation.\cite{142,147} Ti-MOF showed an absorption edge of 350 nm, while the edge of Ti-MOF-NH$_2$ reached ~500 nm. In the photocatalytic process, the organic linker absorbed visible light, and the electrons transferred from its excited state to the CB of the photoactive titanium-oxo cluster, as shown in Figure 9. Pt/ Ti-MOF-NH$_2$ exhibited efficient photocatalytic activity for the
HER under visible light irradiation, and a H$_2$ evolution rate of $\approx$367 $\mu$mol h$^{-1}$ g$^{-1}$ was obtained. In contrast, Pt/Ti-MOF displayed no photocatalytic activity for the HER. These results suggested that the band gap/visible light response played a critical role in the photocatalytic water splitting. However, the structural stability of Pt/Ti-MOF-NH$_2$ should be carefully addressed in the future.

Although MOF-based photocatalysts with organic linkers were successfully developed for water splitting under visible light, the longest wavelength available was only 500 nm due to the limited visible light response and low oxidation power of the BDC-NH$_2$ organic linker.$^{[142]}$ Thus, it is critical to extend the wavelength range by using other organic linkers that have wide visible light absorption and high oxidation power. Bis(4-carboxyphenyl)-terpyridine Ru(II) (Ru(tpy)$_2$) was found to have wider absorption in the visible light region and a lower highest occupied molecular orbital (HOMO) level than BDC-NH$_2$. A Ru complex-modified Ti-based MOF (Ti-MOF-Ru(tpy)$_2$) was synthesized and exhibited high photocatalytic activity for the HER under visible light, with a wavelength up to 620 nm.$^{[148]}$

In addition to the reduction in the band gap of MOF, the visible light response can be increased by developing a dye-sensitized system.$^{[44, 149]}$ Since both MOFs and dyes contain a benzene ring, the strong $\pi$-$\pi$ stacking and van der Waals interactions between MOFs and dyes could enhance the electron transfer in a dye-sensitized system.$^{[150]}$ For example, a rhodamine B (RhB)-sensitized MOF was developed for the HER.$^{[143]}$ RhB-sensitized UiO-66 exhibited high activity for HER under visible light irradiation with a $H_2$ evolution rate of 116 $\mu$mol h$^{-1}$ g$^{-1}$, which was 30 times greater than that of Pt@UiO-66.$^{[149]}$ However, the poor stability of RhB restricted its application as an effective photosensitizer.$^{[151]}$ Recently, Erythrosin B (ErB) dye, with superior stability to RhB, was selected to sensitize UiO-66 MOF for photocatalytic $H_2$ production.$^{[144]}$ With the optimized amount of ErB, Pt-Uio-66-30 (ErB/photocatalyst weight ratio of 3) displayed the highest $H_2$ evolution rate of 460 $\mu$mol h$^{-1}$ g$^{-1}$, while Pt-Uio-66 without sensitization showed no HER activity.

In addition, the photocatalytic activity of ErB-sensitized UiO-66 was higher than that of ErB-sensitized P25-TiO$_2$ and SiO$_2$ for the HER, demonstrating the advantage of MOFs as photocatalysts for water splitting. The enhancement in photocatalytic $H_2$ production was attributed to the efficient transfer of the charge carriers from the photoexcited ErB to the MOFs. These results suggested that the dye-sensitized MOFs showed great potential for the photocatalytic HER.

In addition to the dye-sensitized system based on MOF-based photocatalysts, MOF itself can be used as a photosensitizer for PEC water splitting.$^{[152]}$ MIL-125 (a Ti-based MOF) has a more negative lowest unoccupied molecular orbital (LUMO) level than the CB edge of TiO$_2$, making electron transfer from MIL-125 to TiO$_2$ feasible.$^{[153]}$ The performance of TiO$_2$ for PEC water oxidation was significantly enhanced by the addition of a MOF-based photosensitizer under visible light irradiation.$^{[152]}$ The photocurrent of a TiO$_2$ nanowire in PEC water splitting was improved by 100% from 10 to 20 $\mu$A cm$^{-2}$ at 0.75 V vs. RHE under visible light irradiation after the sensitization of aminated Ti-based MOFs on TiO$_2$. Considering the diversity of MOFs, a large number of MOFs are expected to function as photosensitizers for solar-driven energy conversion. In another study, functionalized MOF-253-Pt was prepared by immobilizing a Pt complex in microporous MOF-253 using a post-synthesis method. MOF-253-Pt acted as both a photosensitizer and a photocatalyst for visible-light-driven HER.$^{[154]}$ The photocatalytic activity of MOF-253-Pt was $\approx$5 times greater than that of the Pt complex, while MOF-253 displayed almost no activity for the HER. Further studies should be conducted to design a robust and highly active MOF bifunctional photocatalyst/photosensitizer for visible-light-driven HER.

3.2.2. Active Site/Co-Catalyst Selection, Incorporation and Optimization

Numerous efforts have been made for the integration of different molecular functional components and MOFs to enhance water splitting activity.$^{[83–87]}$ The combination of MOFs with functional inorganic materials, such as metal NPs and metal oxides, has been reported as a new strategy and has received increased attention.$^{[87, 155, 156]}$ Some researchers incorporated Pt NPs into MOFs to enhance the water splitting activity.$^{[87, 155, 156]}$ The incorporation of Pt NPs into NH$_2$-MIL-101(Cr) was effective for visible-light-driven $H_2$ evolution with high activity.$^{[155]}$ Pt NPs with an average size of $\approx$3.75 nm were highly dispersed on NH$_2$-MIL-101(Cr) with a particle size of $\approx$50 nm. When the Pt amount was increased from 0.5 to 1.5 wt.%, the photocatalytic activity of Pt/NH$_2$-MIL-101(Cr) was improved due to the increased active sites, and the optimum Pt loading for $H_2$ evolution was 1.5 wt.%. A further increase in the Pt amount to 3.0 wt.% decreased the activity due to the agglomeration of Pt NPs. Similarly, Pt NPs with a particle size of 1 nm were embedded into layers of a titanium picolinate framework (TiPF) by photodeposition.$^{[156]}$ The obtained Pt-loaded TiPF could be used as a photocatalyst for the HER, with a hydrogen evolution
rate of 1593 µmol h⁻¹ g⁻¹. Furthermore, TiPF, without Pt NPs, exhibited much lower activity for the HER, with an evolution rate of 228 µmol h⁻¹ g⁻¹, suggesting that Pt active sites played a critical role as an electron trap to improve the charge separation efficiency and increase the HER activity. However, the stability of the TiPF support must be improved in future research.

Recently, earth-abundant Co and Ni-based catalysts have been used to replace the precious Pt metal in water splitting.\[157,158\] Compared to Pt, the control and reduction of the particle size of the oxides are more difficult. Han et al. utilized an MIL-101 MOF to immobilize Co₂O₄ NPs for the OER.\[157\] With cobalt loading in the range of 1.4–4.9 wt.%, ultra-small Co₂O₄ NPs (2–3 nm) were successfully immobilized and well dispersed in the cages of MIL-101. This photocatalyst displayed enhanced photocatalytic activity for the OER, with a high TOF of 0.012 s⁻¹ per Co atom, which was more than 9 times higher than that of the unsupported Co₂O₄ NPs. Furthermore, MIL-101 also promoted the charge transfer process in the OER.

Due to their large BET surface areas, most available MOF and NPs-modified MOF photocatalysts are highly dispersed in water, making them difficult to recycle. It is therefore critical to develop MOF-based photocatalysts that can be easily collected for reuse. One solution is to incorporate magnetic materials into MOFs.\[159–162\] For example, a photocatalyst composed of MIL-53(Fe) microrods with magnetic and active Fe₂O₃ nanosphere decoration on the surface (MIL-53(Fe) hybrid magnetic composites, MHMCs) was reported.\[161\] As shown in Figure 10, the MHMCs were composed of well-dispersed Fe₂O₃ nanospheres with a particle size of ≈200 nm attached to the surface of MIL-53(Fe) microrods, creating the heterostructures. Used as a photoanode, the MHMCs delivered a remarkable photocurrent under visible light irradiation. After the OER, the used MHMCs were easily collected through magnetic separation and were recycled for further use. This structural design approach can provide an ideal platform for the development of MOF-based materials with high activity and magnetic recyclability for energy conversion and environmental remediation.

### 3.2.3. Structural Evolution, Coupling and Creation of Heterojunctions

The charge transfer/separation efficiency is vital to the achievement of high photocatalytic activity. Several strategies have been employed to enhance the charge transfer/separation efficiency, such as developing new MOF photocatalysts through structural evolution and coupling with carbon-based materials to form hybrid photocatalysts.\[162–169\] For example, a highly active Bi-based MOF photocatalyst composed of Bi⁴⁺ and H₃mna (2-mercaptonicotinic acid) was developed (denoted as Bi-mna) for water splitting under visible light.\[162\] According to the experimental and theoretical results, a ligand-to-ligand charge transfer (LLCT) process was responsible for the high photocatalytic activity, which resulted in a longer lifetime of the photogenerated charges, suppressed electron-hole recombination and increased the photocatalytic activity. These results suggested that Bi-based MOFs are promising candidates for efficient water splitting.

In addition to the design of new photocatalysts, functional additives have been used to enhance the charge transfer capability of MOF-based photocatalysts.\[163\] A new hybrid photocatalyst created by coupling UiO-66 octahedrons and g-C₃N₄ (referred to as UG-x, where x is the weight of g-C₃N₄ in the hybrids) was demonstrated.\[163\] Although both g-C₃N₄ and UiO-66 were capable of the HER, the use of each material alone displayed limited photocatalytic activity due to the high charge recombination rate and poor sunlight absorption capability. Figure 11a shows a schematic illustration of the coating of g-C₃N₄ on UiO-66 octahedrons by annealing. UiO-66 had very smooth and clean surfaces as shown in Figure 11b. After annealing, g-C₃N₄ nanosheets were coated on the surface of UiO-66, making the surface uneven (Figure 11c). Increased g-C₃N₄ resulted in heavy coverage of g-C₃N₄ on UiO-66 (Figure 11d). At the optimum weight of g-C₃N₄ in the hybrid, the UG-50 photocatalyst showed dramatically enhanced photocatalytic activity for the HER, with a H₂ evolution rate of 1410 µmol h⁻¹ g⁻¹, compared with UiO-66 and g-C₃N₄ (0 and 80 µmol h⁻¹ g⁻¹, respectively). These results confirmed that the annealing process promoted the formation of UiO-66/g-C₃N₄ heterojunctions and enhanced the charge separation efficiency in UG-50, which enhanced the H₂ evolution rate (more than 17-fold). These results revealed the essential role of the UiO-66/g-C₃N₄ interfaces in enhancing the photocatalytic H₂ production.

Loading CdS with RGO is a useful method to reduce the possibility of electron-hole pair recombination since graphene has high conductivity and superior electron mobility, which can accelerate the electron transfer in photocatalytic reaction.\[166\] Lin et al. found that the hydrogen production performance of CdS was improved by the combined action of UiO-66 and RGO due to the increased quantity of active sites and the minimized recombination of charge carriers.\[166\] As a support, UiO-66 was superior to inorganic semiconductor material. The UiO-66/CdS/1% RGO photocatalyst displayed a very high photocatalytic HER rate, which was 13.8, 2 and 1.23 times greater than CdS, P25/CdS/1% RGO and UiO-66/CdS, respectively, as shown in Figure 12.

In another study, highly porous Co₂O₃@carbon derived from ZIF-67 MOFs was successfully embedded on the surfaces of BiVO₄ nanosheets,\[89\] in which the construction of a p-n heterojunction between the porous Co₂O₃ (p-Co₂O₄) and BiVO₄ significantly promoted the charge transfer and reduced electron-hole recombination. The p-Co₂O₃@carbon/BiVO₄ photoanode demonstrated ≈10 times higher photocurrent density than the bare BiVO₄ and bulk Co₂O₃/BiVO₄ for the OER. This system also demonstrated good stability for the OER without noticeable degradation after three recycled uses. More recently, a highly active Au@CdS/MIL-101 heterostructure was successfully prepared.\[165\] The MIL-101(Cr) with large specific surface area was utilized as a support for well-dispersed Au NPs, and CdS was selectively coated on the Au NPs.\[165\] The good dispersion of CdS and Au NPs, as well as the strong surface plasmon resonance absorption of Au, led to enhanced electron separation and transfer. Under visible light irradiation, the Au@CdS/MIL-101 heterostructure presented a high H₂ production rate of 25,000 µmol h⁻¹ g⁻¹, which was 3.6 and 1.3 times greater than those of CdS and...
CdS/MIL-101, respectively. These results suggested that the heterojunction in the photocatalysts played a critical role in the high photocatalytic activity, and these studies will encourage extensive research on heterostructured photocatalysts in the future.

Besides the separation efficiency of the photo-induced charge carriers, the photocatalytic activity of the MOF-based/derived photosensitizer/photocatalyst essentially relies on the excited state lifetime of the photo-generated electron-hole pair.[45,87,167–173] For example, Lin and co-workers synthesized a MOF based on \{M[4,4′-dcbpy]2bpy\}_2^{2+} building blocks (where M = Ru or Os, bpy = 2,2′-bipyridine and dcbpy = dicarboxy-2,2′-bipyridine), which can be readily excited to their long-lived triplet metal-to-ligand charge transfer (3MLCT) states.[171] The lifetimes of Ru_{2+} excited states in the Os-doped MOF decreased progressively with increasing Os doping amount from 0.3 to 2.6 mol.%. The lifetime at 620 nm decreased from 171 ns in the pure Ru-MOF to 29 ns in the sample with 2.6 mol.% Os doping. In the mixed-metal samples, energy transfer was observed with an initial growth in Os emission corresponding with the decay rate of the Ru excited state. These results demonstrate rapid, efficient energy migration and long distance transfer in these isomorphous MOFs.

Morris and co-workers have made outstanding contributions to the study of the lifetimes of the photo-induced electron and holes in MOF-based photocatalyst.[174–177] The effect of the doping amount of ruthenium(II) tris(5,5′-dicarboxy-2,2′-bipyridine), Rudcbpy, on the lifetime of the emissive 3MLCT state of the Rudcbpy-doped UiO-67 MOF (Zr_{6}(µ_3-O)_{4}(µ_3-OH)_{4}(bpdc)$_6$ (bpdc = biphenyldicarboxylic acid)) was systematically investigated by Morris and co-workers.[174] They found that the lifetime of the emissive 3MLCT state corresponding to Rudcbpy centers incorporated into the MOF backbone was found to be sensitive to the Rudcbpy doping amount in the material. The excited state properties of the Rudcbpy-doped UiO-67 at low doping concentrations resembled that of Rudcbpy in

Figure 10. SEM images of Fe$_3$O$_4$ (a and b), MIL–53(Fe) (c and d) and MHMCs (e and f). Reproduced with permission.[161] Copyright 2015, Royal Society of Chemistry.
dimethylformamide (DMF) displaying a long-lived (≈1.4 µs) \(^{3}\text{MLCT}\) state. Increasing the Rudcbpy doping concentration in UiO-67 was accompanied by a significant decrease in the emission lifetime, which was proposed to be due to the homogeneous energy transfer between the Rudcbpy centers.\(^{[174,175]}\)

Single Rudcbpy preferentially occupied the larger octahedral cages of UiO-67 by incorporation into the backbone of the cage and experienced a DMF-like solvation environment. At higher doping concentrations of Rudcbpy, in addition to incorporation of Rudcbpy into the backbone of the octahedral cavities, populations of encapsulated Rudcbpy were also found in separate octahedral UiO-67 cavities. The decreased lifetime of the slow phase with increased doping concentration was attributed to the intermolecular energy transfer between neighboring Rudcbpy molecules incorporated into the backbone of the octahedral cages.

In another study, Morris and co-workers found that this Rudcbpy-doped UiO-67 can be grown onto conductive fluorine-doped tin oxide (FTO) coated glass substrates without changing its excited state properties or dynamics.\(^{[175]}\)

In addition, it was found that the coupling of the nanosized carbon nitride nanosheets (CNNS) with UiO-66 via a facile electrostatic self-assembly method increased the lifetime of the photo-generated electron-hole pair of CNNS photocatalyst.\(^{[178]}\)

3.2.4. A Brief Summary

The catalytic activities of some typical MOF-based and MOF-derived photocatalysts for the HER and OER are listed in Table 2. In addition to the main active components, the activity of the photocatalysts for the HER and OER is heavily dependent on the sacrificial reagent and the co-catalyst/active sites. Compared to MOF-based electrocatalysts, in which the MOFs were more frequently used as precursors, MOFs have been widely used as photocatalysts directly, although they were often coupled with active sites/co-catalysts to enhance their activity. The amine-functionalized Ti-MOF displayed superior photocatalytic activity for the HER due to the improved visible light response. The addition of metal/metal oxide improved the photocatalytic activity. However, the addition of an excessive amount of metal/metal oxide reduced the photoactivity for the HER and OER. For Pt, a loading amount of 1.5 wt.% in the Ti-MOF-NH2 photocatalyst resulted in the highest \(\text{H}_2\) evolution rate. Similarly, for CoOx NP-modified MOF, the optimum CoOx amount of 3.9 wt.% was demonstrated for the OER, suggesting that the amount and the particle size of the co-catalyst should be controlled and optimized to achieve high photocatalytic activity for the HER and OER. In addition to metals and metal oxides, functional materials, such as CdS, RGO, MoS2 and g-C\(_3\)N\(_4\), have been added to MOF to improve the photocatalytic activity for the HER. As shown in Table 2, MoS\(_2\) increased the photocatalytic activity for the HER and was superior to Pt as a co-catalyst.\(^{[130]}\) The \(\text{H}_2\) production rate of UiO-66/CdS increased 25-fold when
1.5 wt.% MoS₂ was incorporated. The construction of dye-sensi-
tized systems can contribute to high photocatalytic activity; the
selection of the dye, its amount and the incorporation method
should be carefully investigated. In addition to the direct use of
MOFs as photocatalysts, nanostructures and composites derived
from MOFs were investigated as efficient photocatalysts for the
HER and OER. Porous Co₃O₄ nanocages derived from PBA
Co₃[Co(CN)₆]₂ displayed high activity for the OER. However,
the recycled performance of many MOF-based photocatalysts
must be enhanced. Additionally, in most of the studies summ-
arized in this review, the quantum yield of the MOF-based
photocatalysts for water splitting was not presented, and the
values that were presented are rather low and must be improved
in the future.

### 4. Challenges and Perspectives

In this review, the recent research progress of MOF-based and
MOF-derived catalysts for electrocatalytic and photocatalytic
water splitting reactions has been summarized. The applica-
tions of MOFs as direct catalysts and as precursors of catalysts
for water splitting were included, and both the OER and HER
were considered. MOFs have demonstrated promising appli-
cations in the field of electrocatalytic and photocatalytic water
splitting for H₂ production, and several impressive results are
available. However, the field of MOF-mediated water splitting
is in its infancy, and remarkable enhancements are needed to
make MOF-mediated water splitting systems fully competitive.
For example, the poor stability of MOF-based catalysts in water

### Table 2. Photocatalytic activity of some typical MOF-based and MOF-derived photocatalysts for the HER and OER.

| No. | Photocatalyst                          | Target reaction | Sacrificial reagent | Production rate (µmol h⁻¹ g⁻¹) | Quantum yields (%) | Recycled times | Ref. |
|-----|---------------------------------------|----------------|---------------------|-------------------------------|--------------------|----------------|------|
| 1   | Bi-based MOF                          | OER            | AgNO₃               | ≈180                          | /                  | /              | [162]|
| 2   | Ti-MOF                               | HER            | Triethanolamine (TEOA) | 0                             | /                  | /              | [142]|
| 3   | Ti-MOF-NH₂                            |                |                     |                               |                    | /              |      |
| 4   | 0.5 wt.% Pt/Ti-MOF-NH₂                |                |                     |                               |                    | /              |    |
| 5   | 1.5 wt.% Pt/Ti-MOF-NH₂                |                |                     | ≈500                          | ≈1.3 at 420 nm     | 3              |      |
| 6   | 2 wt.% Pt/Ti-MOF-NH₂                  |                |                     | ≈460                          | /                  | /              |      |
| 7   | 2.6 wt.% CoOx NPs-MIL-101             | OER            | [Ru(bpy)₃]²⁺–Na₂S₂O₈ | ≈11,000                       | /                  | /              | [157]|
| 8   | 3.9 wt.% CoOx NPs-MIL-101             |                |                     | ≈15,000                       | /                  | 2              |      |
| 9   | 4.9 wt.% CoOx NPs-MIL-101             |                |                     | ≈13,000                       | /                  | /              |      |
| 10  | Small-sized Ni NPs anchored in MOF-5   | HER            | TEOA                | 3,022                         | 7.8 at 520 nm      | 4              | [158]|
| 11  | MnTD⊂MIL-101                          | OER            | Ceric ammonium nitrate | 2,250                         | /                  | /              | [84] |
| 12  | MIL-101                               |                |                     | 125                           | /                  | /              |      |
| 13  | UIO-66                                | HER            | Na₂S₃, Na₂SO₃       | 0                             | /                  | /              |      |
| 14  | UIO-66/CdS                           |                |                     | 1,700                         | /                  | 6              |      |
| 15  | UIO-66/CdS/1%RGO                      |                |                     | 2,100                         | /                  | /              |      |
| 16  | MoS₂/UiO-66/CdS                      | HER            | Lactic acid         | 32,500                        | 23.6 at 420 nm     | 4              | [143]|
| 17  | UIO-66/CdS                           |                |                     | 1,250                         | /                  | /              |      |
| 18  | g-C₃N₄                               | HER            | L-ascorbic acid     | 80                            | /                  | /              |      |
| 19  | UIO-66                               |                |                     | 0                             | /                  | /              |      |
| 20  | g-C₃N₄/UiO-66 (1:1, w/w)              | HER            | TEOA                | 1,141                         | /                  | /              |      |
| 21  | Ti-MOF-Ru(tpy)₂                      | HER            | TEOA                | ≈200                          | 0.2 at 500 nm      | 3              | [148]|
| 22  | Pt complex immobilized MOF-253       | HER            | CH₃CN               | ≈58,000                       | 1.63 at 440 nm     | 2              | [154]|
| 23  | Pt@UiO-66                            | HER            | TEOA                | 3.9                           | /                  | /              | [149]|
| 24  | 2.54 mg g⁻¹ of RhB@Pt@UiO-66         | HER            |                     | 5.6                           | /                  | /              |      |
| 25  | 11.92 mg g⁻¹ of RhB@Pt@UiO-66        | HER            |                     | ≈100                          | /                  | 3              |      |
| 26  | Erb dye-sensitized Pt/UiO-66          | HER            | Methanol            | 460                           | 0.25 at 420 nm     | 3              | [44] |
| 27  | Hollow Fe₂O₃-TiO₂-PdO₈                | HER            | Lactic acid         | 1,100                         | /                  | 5              | [145]|
| 28  | Porous Co₃O₄ nanocages                | OER            | [Ru(bpy)₃]²⁺–Na₂S₂O₈ | ≈7,900                        | /                  | /              | [103]|
| 29  | Porous Mn₉Co₃O₈ nanocages              | OER            |                     | ≈4,900                        | /                  | /              |      |
| 30  | Porous Fe₃Co₃O₈ nanocages             | OER            |                     | ≈4,000                        | /                  | /              |      |
| 31  | Co₃O₄/TiO₂ p–n heterojunction        | HER            | Methanol            | 7,000                         | /                  | /              | [88] |
is a major concern, which is attributed to the higher affinities of their ions for water molecules than the carboxylate ligands. The leaching of metallic and organic components from the MOFs into the reaction medium during the water splitting reactions is another major concern. The reaction mechanisms of MOF-based catalysts for water splitting are still not well understood and require further investigation. The recycling of MOF-based catalysts is also a major challenge due to the ultrafine size of most MOF-based materials. The main challenges and perspectives for MOF-based catalysts in electrocatalytic and photocatalytic water splitting reactions are discussed in the following sections.

4.1. MOF-Based Catalysts for Electrocatalytic Water Splitting

There are three strategies to design MOF-based electrocatalysts for electrocatalytic water splitting: morphology control, nanostructuring, constructing hybrids/composites and functional modifications. The morphology of the catalyst plays an important role in achieving high electrocatalytic activity. Electrocatalysts with special morphologies, such as CPHs and nanoctahedrons, may show superior electrocatalytic activity than their related NPs. By tailoring the composition of the MOFs and the calcination temperature and atmosphere, various morphologies can be obtained with different catalytic properties for the water splitting reactions. For example, a MOF with a flower-shaped morphology was developed with high electrocatalytic activity. However, due to the diversity of MOFs, it is hard to identify the most efficient morphology for the electrocatalytic OER/HER. A more systematic experimental investigation or theoretical study of the morphology-activity relationship of MOFs is required.

In addition to the morphology, the interaction or synergistic effect between components in the electrocatalysts has a strong influence on the electrocatalytic activity of MOF-derived materials and requires more attention in the development of MOF-based and MOF-derived electrocatalysts for the water splitting reactions. For MOF-derived metal and metal oxide-b catalysts, the leaching of metals during the operation is a major concern and could cause a serious decrease in the electrocatalytic activity and stability for water splitting in acidic or alkaline electrolytes. An effective solution to the deterioration is to encapsulate metal NPs into carbon nanotubes or carbon shells. The use of MOFs as precursors for the synthesis of core-shell structured hybrid electrocatalysts may provide a new research direction for the development of MOF-based catalysts for efficient water splitting. Despite the various MOF-derived nanostructures, precise control of the morphology of these materials is absent because of the limited knowledge of the transformation process. A clearer understanding of this issue is urgently needed and would be helpful for the design and construction of fine nanostructures from MOFs with high surface areas, regular pores and tunable compositions for the electrochemical water splitting reactions.

Imidazolate linkers are often used as functional modifiers to improve the proton transfer process in the electrocatalytic water splitting reactions, which may be applied as a general method to design MOFs with improved charge transfer capability. In addition, functional carbon-based materials, such as GO and rGO, can be used as modifiers for MOF-based electrocatalysts to improve the charge transfer process. For example, CoP/rGO can be used as a bifunctional catalyst on both the anode and cathode for water splitting in alkaline solutions, with better activity (135 mV dec⁻¹) than the integrated Pt/C and IrO₂ catalyst couple (201 mV dec⁻¹).[129] However, the mechanism must be clarified. In addition, other carbon-based materials, such as C₃N₄, could be used to improve the charge transfer efficiency in the electrocatalytic OER and HER.

4.2. MOF-Based Catalysts for Photocatalytic Water Splitting

As a photocatalyst for water splitting, the efficiency of a MOF-based material is strongly affected by its sunlight absorption capability (band gap), the number of active sites and the charge separation/transfer efficiency. This review provides a summary of recent developments involving the use of MOFs as photocatalysts, with particular emphasis on several key strategies (band gap engineering, photosensitization, active site/co-catalyst selection and optimization and the coupling of MOFs with other functional materials) to develop MOF-based and MOF-derived photocatalysts with high performance.

The band gap energy of MOFs can be tailored by changing the metal-oxo clusters and bridging organic linkers. Several organic dyes have been used to sensitize MOF-based photocatalysts, leading to significantly enhanced photoactivity under visible light irradiation. However, the stability of these organic-linker-modified MOFs and the organic dyes should be carefully considered in future research. MOFs themselves can also be used as photosensitizers for PEC water splitting and can significantly enhance the photoactivity of TiO₂ under visible light.[139] In the future, the match of the band structure of MOFs and TiO₂ should be clarified by theoretical calculations, such as DFT, which may provide useful design guidelines for MOF-based photosensitizers.

The incorporation of active sites, such as Pt NPs, into MOFs was found to dramatically enhance the photocatalytic activity.[142,143] However, the limited reserves and high cost of Pt hinder its large-scale application. Recently, highly efficient, small-sized Ni particles embedded in MOF-5 as co-catalysts/active sites with low overpotential for photocatalytic HER under visible light was demonstrated.[145] A low overpotential of ~0.37 V was obtained by this photocatalyst, which was comparable to that of Pt@MOF-5. The development of non-precious metal NPs in MOFs to enhance the photocatalytic activity requires more attention in the future. It is critical to develop MOF-based photocatalysts that can be easily collected and reused. One solution is to incorporate magnetic materials into the MOFs. After water splitting, the magnetic photocatalyst can be easily collected by magnetic separation and recycled for further use. Magnetic MOF-based composites should be next-generation photocatalysts for water splitting.

The charge transfer/separation efficiency plays a vital role in the photocatalytic water splitting reactions. Various methods have been used to improve the charge transfer/separation efficiency. For example, electrons can be transferred from excited organic linkers to metal-clusters in Zr-based MOFs (LLCT
process). In the H₂ generation process via the LLCT mechanism, the CB edge position of a titanium-oxo cluster is more suitable for charge transfer since the CB potential of a titanium-oxo cluster is more positive than that of its zincinum counterpart. Thus, Ti-based MOFs are expected to be highly active photocatalysts for the HER. In addition to the design of new photocatalysts, functional additives, such as g-C₃N₄, have been used to enhance the charge transfer/equilibrium efficiency of MOF-based photocatalysts. For example, g-C₃N₄/UiO-66 hybrid photocatalysts showed enhanced activity in the visible-light-driven HER, which was attributed to the efficient interfacial charge transfer from photoexcited g-C₃N₄ to UiO-66. These results demonstrated the potential use of MOFs and g-C₃N₄ to construct active photocatalysts for water splitting. Based on the extensive studies of the application of MOFs in photocatalytic water splitting, the application of MOFs in solar cells will be a research focus in the future, although it is out of the scope of this article. It will be exciting to watch the rapid development of such new materials in the years to come.

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