

## REVIEW

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## Photocatalytic hydrogen production: an overview of new advances in structural tuning strategies

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Hydrogen production from photocatalytic water decomposition provides a promising approach to harvest solar energy for addressing the challenges of energy shortage and environmental pollution. To meet the commercialization target set by the US Department of Energy, three great challenges of photocatalytic hydrogen production should be well addressed: the capture of solar light, the separation/transfer of photo-excited charge pairs and the surface redox reactions to produce useable fuels. The component, morphological and structural designs are highly promising solutions for solving these challenges. Among them, structural designs are more effective and easier; in particular, the incorporation of single atoms into the structure, defect control and formation of S-scheme heterojunctions has attracted considerable attention. The recent advancements in these topics are highlighted and discussed in this review.

## Introduction

Energy shortage and environmental pollution are two major issues hindering the sustainability of modern civilization.<sup>1–3</sup> The search for sustainable and green energy sources is extremely urgent and highly demandable. Among all the intensively studied alternative energy sources, such as wind and tidal power, solar energy is an energy source that can meet our long-term needs because of its advantages of zero cost and being abundant and inexhaustible.<sup>4–8</sup> Recently, conversion of solar energy to useable fuels (*e.g.*, hydrogen, ethane, and propanol) using semiconductor photocatalysts has attracted considerable attention,<sup>9–12</sup> especially in the fields of solar-driven overall water splitting, photocatalytic CO<sub>2</sub> reduction and

photocatalytic degradation of pollutants.<sup>13–17</sup> Among the useable fuels, hydrogen is a highly efficient energy source because of its ultra-high energy density and zero emission.<sup>18,19</sup> Photocatalytic water decomposition provides a promising approach to harvest solar energy for generating environment-friendly hydrogen energy for our long-term needs.<sup>20–22</sup> The hydrogen production cost from solar-driven water splitting should be limited below \$2.0 kg<sup>−1</sup> on the terawatt scale to compete with the traditional hydrogen production methods, such as coal gasification, in the global market. To achieve this goal, analysis suggests a solar-to-hydrogen efficiency (STH) of 10% as a threshold for energy return on investment.<sup>23</sup>

Photocatalytic processes involve three essential steps: the capture of solar light, the separation/transfer of photo-excited charge pairs, and the surface redox reactions to produce useable fuels.<sup>23–26</sup> To reach such STH, challenges in the photocatalytic processes should be addressed. Additionally,

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a photocatalyst should also have decent stability for practical usage. Unfortunately, none of the existing single photocatalysts show decent efficiency in all requirements. For instance,  $\text{TiO}_2$  has a band gap of 3.2 eV, which means that it can only absorb parts of ultra-violet (UV) light. UV light only accounts for less than 5% of the total solar energy, while visible light and infrared light contribute 43% and 52% of solar energy, respectively.<sup>27–29</sup> A good photocatalyst, therefore, should show decent light absorption in the visible light and near infrared light regions. Another example is  $\text{Fe}_2\text{O}_3$  that has a bandgap of 2.1 eV and can absorb visible light up to 590 nm. Unfortunately, it suffers from the fast recombination of photoexcited charge pairs and the average charge carrier migration length is less than 10 nm.<sup>30</sup>

To improve the solar to hydrogen efficiency, many strategies have been proposed. Among them, the search for new materials is the most effective way to improve the STH. Solar-driven hydrogen production began with the observation of water photooxidation on semiconducting  $\text{TiO}_2$ ; its limited STH efficiency, however, is only 1.3% due to a wide bandgap of 3.2 eV.<sup>31</sup> The investigations on semiconducting transition metal oxides with empty d bands (*i.e.*,  $\text{V}^{5+}$ ,  $\text{Nb}^{5+}$  and  $\text{W}^{6+}$ ) show that their valence-band energy levels are dictated strongly by the O 2p levels, making their valence band maximum (VBM) higher than that of  $\text{TiO}_2$  but still thermodynamically capable for water oxidation. In addition, incorporating  $s^2$  cations (*e.g.*,  $\text{Sn}^{2+}$  and  $\text{Bi}^{3+}$ ) can narrow the band gap through a coupling between s and O 2p bands that pushes the VBM upward and another coupling between the empty d bands and the p bands of the  $s^2$  cations that pushes the conduction band minimum (CBM) downward.<sup>32</sup> For instance, our previous work demonstrated that the composition tuning of Bi element, which has both  $d^0$  band ( $\text{Bi}^{5+}$ ) and  $6s^2$  cations ( $\text{Bi}^{3+}$ ), in a  $\text{Ba}_2\text{Bi}_{1-x}\text{Nb}_{1-x}\text{O}_6$ -type double perovskite oxide can greatly shift down the CBM and shift up the VBM and therefore improve the light absorption.<sup>9,10</sup> Novel photocatalysts, such as  $\text{BiVO}_4$ ,  $\text{WO}_3$  and  $\text{Fe}_2\text{O}_3$ , which possess much higher theoretical STH, have been proposed. For instance, the theoretical STH of  $\text{BiVO}_4$  approaches 9.2% with a photocurrent of  $7.5 \text{ mA cm}^{-2}$  under the standard AM1.5G solar light illumination.<sup>33</sup> Accordingly, these materials have garnered increasing attention for decades. Unfortunately, high ionic character of these oxides induces a large separation between the band edges, limiting efficient solar energy harvesting especially in visible and near infrared light regions. Until now, no single semiconductor can achieve the targeted STH because of tremendous difficulties in searching for new materials that can fulfill all three requirements. The most promising materials, therefore, are still the semiconductors that are intensively studied for decades.

Another approach is the structural tuning strategy, which includes the modification (*e.g.*, heteroatom doping and defect control) of existing semiconductors and the fabrication of tandem devices for taking full advantage of synergistic effects among varied semiconductors. Also, the latter has already shown high potential for practical applications with high STH. For instance, in 2017, a photoelectrochemical (PEC) STH of 16.2% was achieved using a Si/GaInAs/GaInP tandem device with a newly found semiconductor, 1.8 eV-bandgap p-type

GaInP.<sup>23</sup> In 2018, the efficiency was improved to 19% using a tandem cell comprising  $\text{TiO}_2$ ,  $\text{AlInPO}_x$ ,  $\text{AlInP}$ ,  $\text{GaInP}$ ,  $\text{GaInAs}$  and  $\text{GaAs}$ , using the same AM1.5 light source.<sup>34</sup> Most recently, the Mi group reported a STH over 9.2% for photocatalytic hydrogen production from pure water and concentrated solar light using  $\text{InGaN/GaN}$  nanowires as the photocatalysts.<sup>35</sup> Besides the three requirements for high conversion efficiency mentioned above, the device must be composed of widely available and low-cost component atoms, be inexpensive to be fabricated and be robust enough for long-term operation. Although these devices manifest high STH, their dramatically high fabrication cost is the greater challenge for hydrogen production cost control below  $\$2.0 \text{ kg}^{-1}$ .

Moreover, the structural tuning strategy is much easier than developing new materials and is more effective than other strategies such as morphology control. For instance, heteroatom doping demonstrates the most effective way to improve the performance of a semiconductor. Doping  $\text{Mo}^{6+}$  or  $\text{W}^{6+}$  into the  $\text{V}^{5+}$  sites of  $\text{BiVO}_4$  increases the electronic conductivity and thus carrier collection efficiencies.<sup>36–39</sup> Incorporation of Cu into  $\text{WO}_3$  induces strong Cu 3d character in the valence band, which raises the VBM and narrows the bandgap.<sup>40,41</sup> The most widely used doping elements are “N” and “S/Se” because the substitution of O by N and S/Se will decrease the bandgap by introducing higher-lying N 2p states and S/Se p-band energy levels, respectively.<sup>42–46</sup> Our recent work shows that the incorporation of S into the perovskite oxide can push the CBM down by S 2p orbital coupling, which pushes the light absorption edge to the near infrared light region. The bandgap tuning of perovskite oxides increases the light absorption but is not at the expense of losing stability due to the natural chemical stability of oxides.<sup>47</sup> Moreover, the fabrication of heterojunction or tandem structures is also one of the effective structural tuning methods. For PEC water splitting, tandem structures already demonstrate the capability for achieving high STH as mentioned above,<sup>23,34,35</sup> while for photocatalytic hydrogen production, a heterojunction formed by the direct contact of two semiconductors is widely used to improve the efficiency of the separation of charge pairs. Three traditional type heterojunctions, *i.e.*, straddling gap, staggered gap, and broken gap, have been intensively studied for decades.<sup>48–50</sup>

It is well-known that the small size of metal catalysts is beneficial to the improvement of metal utilization efficiency, which is particularly important for noble metal components. Classic metal nanoparticles (or nanocrystals) have low metal utilization efficiency due to the existence of abundant bulk atoms. Contrastively, single-atom catalysts (SACs) with a theoretical metal utilization rate of 100% and subnanometer nanoclusters with nearly 100% atom efficiency for specified reactions have received increasing attention in recent years.<sup>51</sup> SACs contain isolated metal atoms or mononuclear metal complexes on appropriate supports, while a nanocluster with well-defined structures in the size regime of a few subnanometers to several nanometers consists of tens to hundreds of metal atoms.<sup>52,53</sup> Both SACs and nanoclusters all have the following advantages over their nanoparticle counterparts: (1) low-coordination environment; (2) quantum size

effects; (3) metal–support interactions, originating from the chemical bonding effect and the charge transfer between metal species and supports.<sup>51</sup> Remarkably, compared to nanoclusters, SACs have exclusively higher activity and selectivity brought by their unsaturated coordination sites and unique electronic structures. In addition, embedding SACs into transition metal oxides has been proved to be an effective strategy not only to minimize the loading amount of precious metals but also to improve the intrinsic performance of catalysts.<sup>54</sup> For example, Sun *et al.* dispersed Ir atoms into the WO<sub>3</sub> matrix lattice *via* a two-step method, which exhibited improved activity and durability for acidic overall water electrolysis.<sup>55</sup> In this case, the valence of an isolated metal single atom should not be necessarily zero to boost the metal–support synergistic effects. More importantly, the appropriate supports can be in the form of clusters, trimers, dimers, monomers, and even nanoclusters. SACs, therefore, could be the ultimate size limit for nanoclusters and broaden the scope of nanoclusters.

To address the irreconcilable conflict between a wide light absorption range and high redox ability in single semiconductors, people have never stopped the studies on heterojunctions since 1984.<sup>56</sup> With the continuous deep understanding on the light absorption and charge transfer in heterojunctions, fine control of the heterojunction structure became more necessary. Recent calculations on PEC tandem devices indicate that theoretical STH can reach 21%, 24% and 27% for a junction with top/bottom bandgaps of 1.7/1.2, 1.8/1.2 and 1.7/1.05 eV, respectively.<sup>23</sup> For a photocatalytic hydrogen production heterojunction, the interpretation of the charge-transfer route is controversial. Recently, the Yu group raised a different charge transfer route (S-scheme heterojunction) compared to the widely acknowledged type-II and Z-scheme heterojunctions.<sup>57</sup>

Given that traditional three type heterojunctions and heteroatom doping have been intensively studied and reviewed previously, in this review, we will brief the fundamentals of structural tuning strategies of the newly emerged approaches, *i.e.*, single atom photocatalysts, defect control and S-scheme heterojunctions. We present an analysis on their advantages for photocatalytic hydrogen production and mainly focus on how these strategies address the three challenges mentioned above in photocatalytic processes. Since no single attempt can solve all challenges, in the final section, we discussed the limitations of each approach, and the future research direction was also discussed.

## Structural tuning approaches

### Single atom metal photocatalysts

To address the issues of rapid recombination of photo-generated electron–hole pairs and insufficient surface active sites on the photocatalyst, SACs have been employed and become a hot topic in water-splitting photocatalysts because of their unique advantages, such as the highest atom utilization rate and excellent catalytic activity.<sup>58–66</sup> Single atoms (SAs) can be incorporated into the crystal lattice of the photocatalyst substrate or in the form of isolated metal atoms or

mononuclear metal complexes on varied substrates. Since Zhang *et al.* firstly reported single Pt atoms on the FeO<sub>x</sub> carrier with a Pt–O–Fe structure in 2011,<sup>67</sup> most atoms in the periodic table can be synthesized as SA materials. Therefore, the versatile nature of SAs and the diversity of photocatalyst substrates can enable the flexible engineering of both electronic structures and crystal structures, thereby facilitating the charge separation, narrowing the bandgap, and increasing the surface reaction kinetics. The material obtained by this method not only has the advantages of a homogeneous catalyst, including the uniformity of active sites and the adjustable interaction with ligands, but also inherits the high durability and excellent recoverability of a heterogeneous catalyst.<sup>68–75</sup> Yang's group prepared single metal atoms (Pt, Pd, Rh, and Ru) on TiO<sub>2</sub> for the first time,<sup>68</sup> and the catalyst showed higher photocatalytic activity than the metal clusters supported on TiO<sub>2</sub> for hydrogen production. This work opened a door for the application of SACs in photocatalysis. The introduction of SACs can not only effectively improve the light absorption capacity, broaden the light collection range, and improve the efficiency of charge separation/transfer, but also maximize the utilization rate of atoms and reduce the use of catalytic metals due to the unique exposure mode. They exhibit remarkable stability and long service life, which has been verified in many photocatalytic reactions.<sup>76–81</sup> Since then, SACs have attracted wide attention and developed rapidly in the field of photocatalysis.

### Noble metal SAs

SAs can not only accept photo-generated electrons and promote the separation and transfer of interface charges, but also provide proton reduction sites.<sup>82</sup> Besides that, noble metal single atoms or nanoparticles are also active photo-absorbers, such as Au. The hot electron flow between noble metal single atoms and the substrate greatly promotes the charge transfer and surface reactions. There are eight kinds of noble metals, which are used in the field of water decomposition photocatalysis. Herein, we will discuss the widely studied four kinds of them, *i.e.*, Pt,<sup>83–86</sup> Rh,<sup>87</sup> Ru,<sup>88</sup> and Ag.<sup>89,90</sup>

**Pt.** Pt is considered to be an ideal cocatalyst because of its proper Fermi energy level to capture electrons and low hydrogen evolution potential.<sup>91,92</sup> Anchoring Pt SAs on the substrates of TiO<sub>2</sub>, C<sub>3</sub>N<sub>4</sub>, and metal organic frameworks (MOF) with large surfaces can optimize the structure of the photocatalyst on the atomic scale, promoting the separation of electron–hole pairs.<sup>93–95</sup> In 2019, Chen *et al.* assembled Pt SAs on a defective TiO<sub>2</sub> support (Pt<sub>1</sub>/def-TiO<sub>2</sub>).<sup>96</sup> Apart from being proton reduction sites, Pt SAs promoted the neighboring TiO<sub>2</sub> units to generate surface oxygen vacancies and formed a Pt–O–Ti<sup>3+</sup> atomic interface. This unique atomic interface effectively facilitated photogenerated electrons to transfer from Ti<sup>3+</sup> defective sites to Pt SAs, thereby enhancing the separation of electron–hole pairs. As a result, the Pt<sub>1</sub>/def-TiO<sub>2</sub> exhibited a record-level photocatalytic hydrogen production performance with an unexpectedly high turnover frequency (TOF) of 51 423 h<sup>−1</sup>, under UV-visible light illumination (300 W Xe lamp). Hu *et al.* (Fig. 1) also investigated Pt SA loaded defective TiO<sub>2</sub> using TiO<sub>2</sub>



Fig. 1 The synthesis strategy for the Pt SA/Def-s-TiO<sub>2</sub> photocatalyst. Copyright 2021, Elsevier.

nanosheets as the substrate.<sup>97</sup> Their computational analysis treated Pt SAs as homogeneous catalysts rather than heterogeneous catalysts in Chen's work. They found that defective TiO<sub>2</sub> nanosheets could effectively stabilize the single-atomic Pt sites through constructing a three-center Ti–Pt–Ti structure. More importantly, the Ti–Pt–Ti structure can hold the stability of isolated single-atomic Pt sites and facilitate the separation and transfer of photoinduced charge carriers, resulting in improvements in photocatalytic H<sub>2</sub> evolution. The photocatalyst delivers a hydrogen production rate of 13 460.71 mol h<sup>−1</sup> g<sup>−1</sup> under UV-visible light illumination (300 W xenon lamp).

Polymeric graphitic carbon nitride (g-C<sub>3</sub>N<sub>4</sub>) has attracted extensive attention in recent years because of its high chemical stability and attractive electronic structure. Li *et al.* took Pt SAs and g-C<sub>3</sub>N<sub>4</sub> as an example to demonstrate that metal–ligand charge transfer could adjust the light absorption and improve the carrier utilization rate,<sup>98</sup> resulting from strong interaction between the d<sup>0</sup> orbitals of Pt and N-forming π-conjugated graphitic planes (Fig. 2). Li *et al.* synthesized a new single Pt atom co-catalyst by anchoring Pt SAs on g-C<sub>3</sub>N<sub>4</sub> with high dispersion and stability.<sup>99</sup> When the Pt loading amount reached 0.38%, Pt atoms dispersed and formed sub-nano clusters. The isolated Pt SAs induced the intrinsic change of the surface trap state, which prolonged the lifetime of photo-generated electrons.

Instability of SAs is a great challenge preventing their massive applications, especially the aggregation issue, since unanchored SAs are too fluid.<sup>51</sup> MOFs or covalent organic frameworks (COFs) are an important class of porous crystals with various customizable structures as well as functions and large surface area. They can provide the most effective coordination to anchor metal SAs, which mitigates the aggregation during the photocatalytic processes. In addition, the interaction between SAs and the MOF/COF substrate produces active sites and further regulates the light absorption capability. For instance, Zuo *et al.* prepared the first SACs based on 2-dimensional (2D) MOFs,<sup>100</sup> as shown in Fig. 3. 12.0 wt% 2D monoatomic catalyst showed a photocatalytic hydrogen production rate of 11 320 mol h<sup>−1</sup> g<sup>−1</sup> under visible light irradiation (λ > 420 nm).

**Ru.** As the cheapest metal in the platinum group, the Ru-based catalyst has proper hydrogen bonding energy, and when it is used as a SAC, it makes a significant contribution to

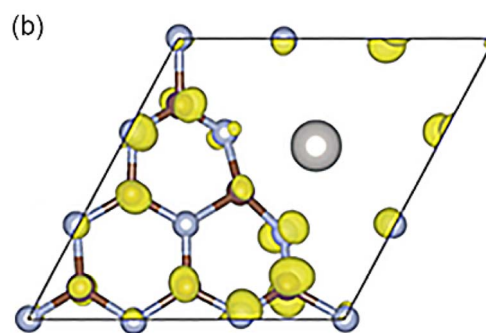
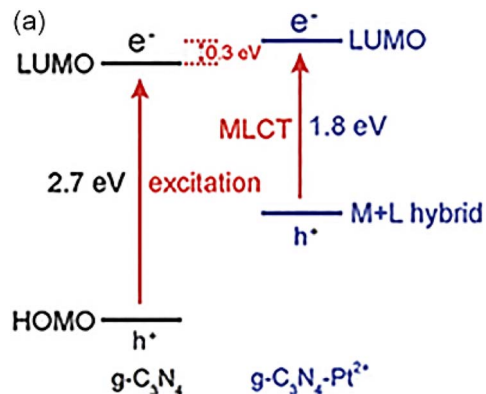


Fig. 2 (a) Energy-band alignment diagrams for g-C<sub>3</sub>N<sub>4</sub> and g-C<sub>3</sub>N<sub>4</sub>-Pt<sup>2+</sup>. (b) Photoexcited charge density transition from the Pt<sup>2+</sup>-induced hybrid highest occupied molecular orbital (HOMO) states to the lowest unoccupied molecular orbital (LUMO) of g-C<sub>3</sub>N<sub>4</sub>-Pt<sup>2+</sup>. Copyright 2016, Wiley-VCH.

the minimum value of the conductive band, thus promoting electron transfer and improving hydrogen adsorption. Zhang *et al.* decorated Ru SAs on the polygonal TiO<sub>2</sub> sphere,<sup>101</sup> and under illumination with a 300 W Xe lamp, its hydrogen evolution rate was 7.2 mmol h<sup>−1</sup> (Fig. 4). Yan *et al.* embedded Ru SAs into N-doped TiO<sub>2</sub>/C carrier (Ru-SAs@N-TC) derived from a MOF of NH<sub>2</sub>-MIL-125.<sup>102</sup> The Ru-free control sample showed negligible photocatalytic activity due to rapid recombination of photoinduced charge carriers. Contrastively, Ru-SAs@N-TC with a Ru loading amount of 0.360 wt% displayed a higher photocatalytic H<sub>2</sub> production rate of 58.3 μmol h<sup>−1</sup> and a TOF of 81.8 h<sup>−1</sup>, under illumination with a 300 W Xe lamp. But their results showed inferior activities for the Ru SA sample to Ru nanoparticles, which was ascribed to the optimal intrinsic activity of Ru nanoparticles. Further work on precisely controlling the atomic coordination environment of Ru SAs should be conducted to further understand the mechanism.

**Pd.** Pd is a cheaper alternative to Pt and usually used as a dopant to modify the activity of photocatalysts for hydrogen production. Given that the efficiency of g-CN, a layered photocatalyst, is still too low due to the poor utilization of photoexcited-charge carriers, Cao *et al.* inserted Pd SAs in the space of adjacent g-CN layers and anchored Pd atoms on the surface of g-CN to synthesize an atom-dispersed Pd layered photocatalyst with adjacent layer bridges and g-CN surface

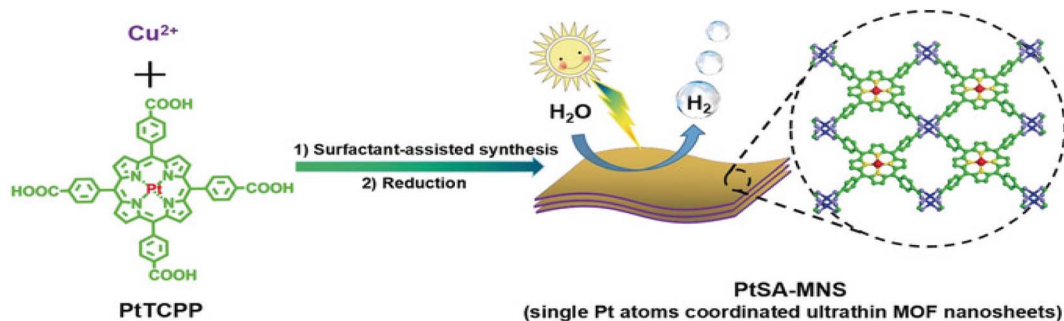


Fig. 3 Illustration of the synthetic route towards Pt single-atom-coordinated ultrathin MOF nanosheets (PtSA-MNSs) for photocatalytic hydrogen production. Copyright 2019, Wiley-VCH.

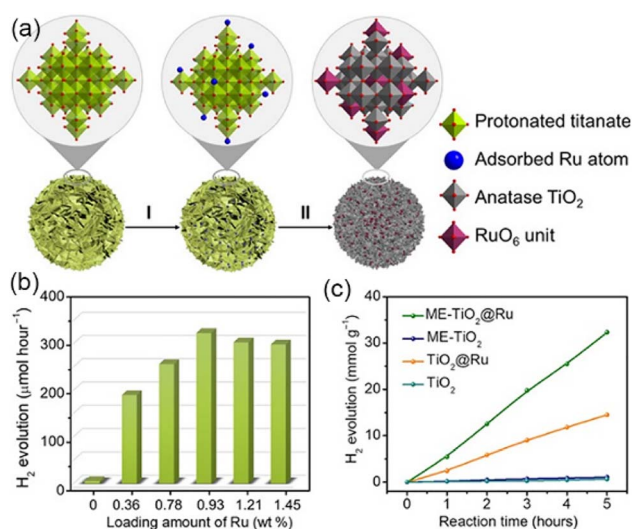


Fig. 4 (a) Illustration of the synthetic route to Ru SAs in TiO<sub>2</sub>. (b) Photocatalytic H<sub>2</sub> generation rates of ME-TiO<sub>2</sub>@Ru with different Ru loading amounts. (c) Time-dependent hydrogen evolution activities. Copyright 2020, the author, and American Association for the Advancement of Science.

sites,<sup>103</sup> as shown in Fig. 5. Both theoretical prediction and empirical characterization confirmed the successful synthesis of a single-atom engineered Pd/g-CN hybrid. The photocatalyst exhibited excellent charge separation and transfer, favoring efficient photocatalytic hydrogen evolution, which is much better than that of the optimized Pt/g-CN benchmark.

**Ag.** Ag has been widely proved to be an excellent photocatalyst, but when it is used as a cocatalyst for H<sub>2</sub> evolution, the hydrogen evolution rate is low due to its lower work function than other noble metals, which limits its photocatalytic performance for water decomposition. Jiang *et al.* prepared an Ag-N<sub>2</sub>C<sub>2</sub>/CN photocatalyst with a new configuration (Fig. 6),<sup>104</sup> high SA loading amount, remarkable activity, and good stability by adjusting the coordination mode of the monatomic metal. The photocatalytic hydrogen evolution activity was significantly higher than that of its nanoparticle counterparts, which was ascribed to expanded light absorption and higher Fermi level position (indicating a higher carrier density and better charge

transfer capability), due to the incorporation of Ag SAs. To address the sluggish surface redox reactions, Li *et al.* fabricated Cu, Ag, Co, and Ni SA embedded g-CN *via* a facile two-step pyrolysis method.<sup>90</sup> The results indicated that Ag SAs displayed the best photocatalytic hydrogen production activities, because Ag SAs were trapped in the six-fold cavity of g-CN and upgraded the  $\Delta G_{H^+}$  of the adjacent pyridinic N atoms rather than directly serving as the active sites.

#### Non-noble metal SAs

Although noble metals are widely used in the field of photocatalytic water decomposition, their scarcity and high price limit the large-scale application, so it is highly demanded to develop efficient and low-cost non-noble metal

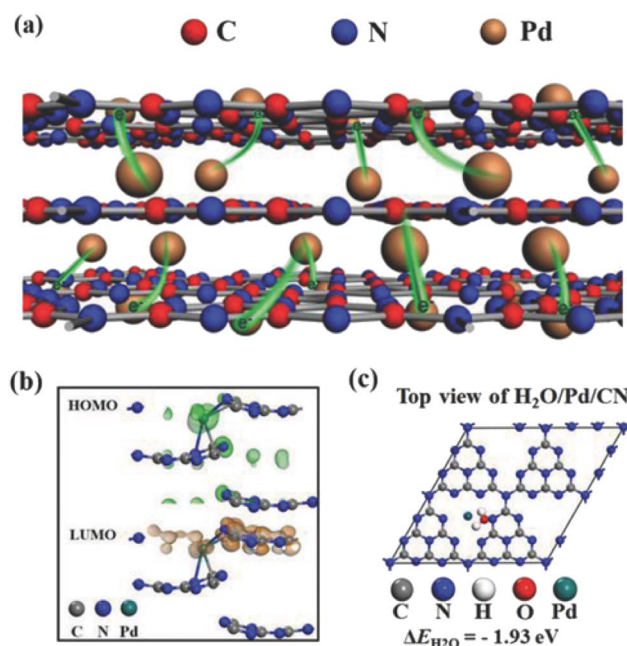


Fig. 5 (a) Conceptual illustration of the Pd/g-CN hybrid with the interlayer intercalation and surface anchoring of Pd atoms. (b) The HOMO/LUMO calculation of Pd/g-CN. (c) The calculation of adsorption energy of a H<sub>2</sub>O molecule on the Pd/g-CN. Copyright 2018, Wiley-VCH.

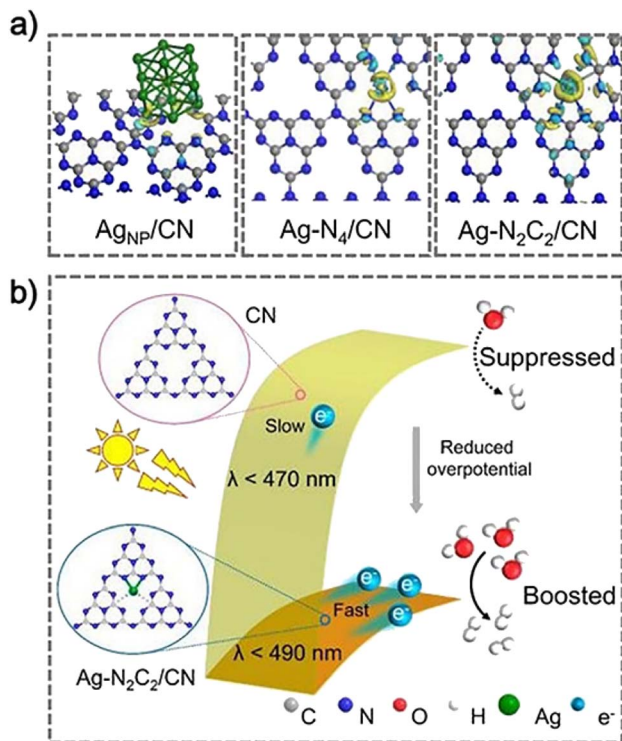


Fig. 6 (a) Charge density difference maps, and (b) proposed mechanism for photocatalytic  $\text{H}_2$  evolution over CN and  $\text{Ag-N}_2\text{C}_2/\text{CN}$ . Copyright 2020, Wiley-VCH.

photocatalysts.<sup>105,106</sup> The introduction of non-noble metal atoms to synthesize photocatalysts not only reduces the utilization rate of noble metals but also breaks the limit of activity noble metal atom sites.

**Co.** As a non-noble metal, the Co atom has attracted more and more attention in the field of photocatalytic water decomposition in recent years. Cao *et al.* proposed a photocatalyst prototype for the first time in 2017,<sup>107</sup> by anchoring Co SAs on carbonitride, and creating an active single  $\text{Co}_1\text{-N}_4$  site on  $\text{g-C}_3\text{N}_4$  for efficient photocatalytic hydrogen production. This noble metal-free composite material showed a high hydrogen production rate of  $10.8 \mu\text{mol h}^{-1}$  under UV-Vis illumination (300 W Xe lamp). To address the challenge of charge separation for robust overall water splitting, a single  $\text{Co}_1\text{-P}_4$  site confined on  $\text{g-C}_3\text{N}_4$  nanosheets was prepared by a facile phosphidation method (Fig. 7).<sup>108</sup> Coordinatively unsaturated Co SA sites could effectively suppress charge recombination and prolong carrier lifetime by about 20 times compared to pristine  $\text{g-C}_3\text{N}_4$ , and boost water molecular adsorption and activation for oxygen evolution. This single-site photocatalyst exhibited steady water splitting activity with  $\text{H}_2$  evolution rate up to  $410.3 \mu\text{mol h}^{-1} \text{g}^{-1}$  under 300 W Xe lamp illumination and a quantum efficiency of 2.2% at 500 nm.

Aiming at mitigating the aggregation or sintering of SACs during the synthetic procedures and to achieve high loading amounts of SACs, Shi *et al.* developed a new *in situ* preparation strategy on carbon nitride,<sup>109</sup> which could synthesize separated cobalt SAs with a high metal loading amount up to 5.9 wt%.

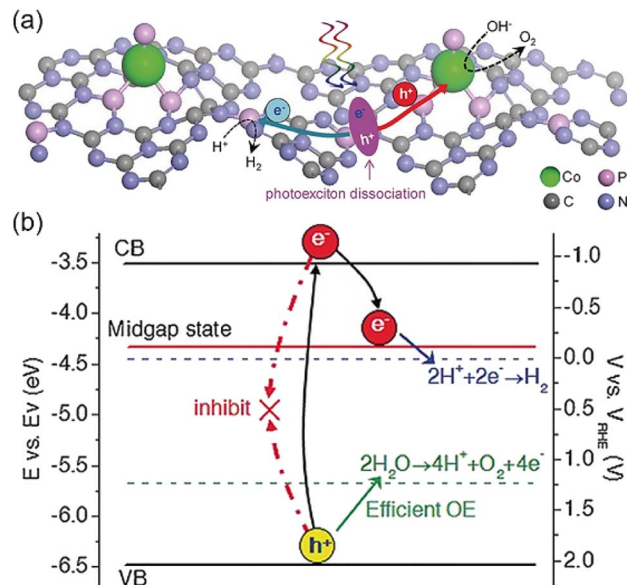


Fig. 7 (a) Schematic illustration of the solar-driven overall water splitting on the  $\text{Co}_1$ -phosphide/PCN photocatalyst, in which the photogenerated electrons and holes ( $\text{h}^+$ ) are spatially separated. (b) Electronic band structure diagram for the  $\text{Co}_1$ -phosphide/PCN photocatalyst. Copyright 2017, Wiley-VCH.

Furthermore, due to the rapid electron transfer from carbon nitride to isolated Co SAs, the photocatalytic  $\text{H}_2$  production rate reached  $1180 \mu\text{mol g}^{-1} \text{h}^{-1}$  using a light-emitting diode light source (12 W,  $\lambda = 420 \pm 10 \text{ nm}$ ).

**Ni.** Xia *et al.* decorated a Ni-imidazole framework on CdS nanorods for photocatalytic hydrogen production.<sup>110</sup> Physicochemical characterization and theoretical calculations revealed strong interactions at the CdS/framework interface and the presence of numerous  $\text{Ni-N}_4$  SA active sites, leading to the significant performance enhancement. Under visible light irradiation ( $\lambda > 420 \text{ nm}$ ), the obtained photocatalyst reached a hydrogen evolution rate of  $21712 \mu\text{mol h}^{-1} \text{g}^{-1}$ , which is 14.23-fold higher than that of pristine CdS. Ni can also be used as the cocatalyst to improve the hydrogen evolution performance. Su *et al.* atomically dispersed Ni in zinc sphalerite cadmium-zinc sulfide quantum dots (ZCS QDs).<sup>111</sup> The photocatalyst delivered an efficient and durable photocatalytic performance for water splitting under visible light irradiation ( $\lambda > 420 \text{ nm}$ ) (Fig. 8). The finely tuned Ni SAs dispersed in ZCS QDs exhibit a photocatalytic  $\text{H}_2$  production activity of  $18.87 \text{ mmol h}^{-1} \text{g}^{-1}$ . The improvements could be attributed to the favorable surface engineering to achieve highly active sites of monovalent Ni(i) and the surface heterojunctions to reinforce the carrier separation owing to the suitable energy band structures, built-in electric field, and optimized surface  $\text{H}_2$  adsorption thermodynamics.

### Defect control

Engineering defects is an effective method to improve the activity of the photocatalyst by adjusting the coordination number and electronic structure of atoms, improving the

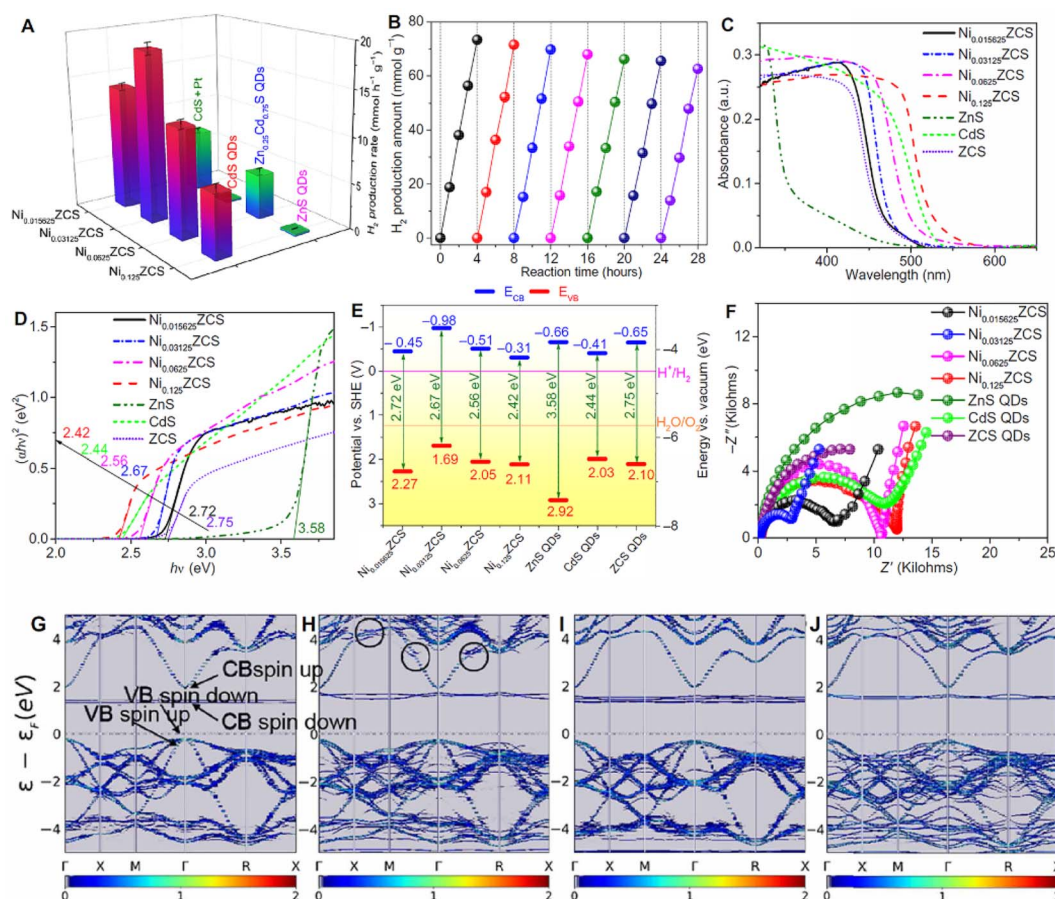


Fig. 8 Photocatalytic performance and spectroscopy/(photo)electrochemical characterization. (A) A comparison of the photocatalytic  $\text{H}_2$  production activities of  $\text{Ni}_x\text{ZCS}$  ( $x = 0.015625, 0.03125, 0.0625, \text{ and } 0.125$ ) QDs, Pt-CdS QDs, CdS QDs, ZCS QDs, and ZnS QDs. The error bars denote SD. (B) Time course of photocatalytic  $\text{H}_2$  production over  $\text{Ni}_{0.03125}\text{ZCS}$ ; every 4 hours, the reaction system was purged with Ar for 30 min to remove  $\text{H}_2$ . (C) Ultraviolet-visible (UV-vis) diffuse reflectance spectra of  $\text{Ni}_x\text{ZCS}$  ( $x = 0.015625, 0.03125, 0.0625, \text{ and } 0.125$ ) QDs, CdS QDs, ZCS QDs, and ZnS QDs. (D) The plots of transformed Kubelka-Munk function of the  $\text{Ni}_x\text{ZCS}$  ( $x = 0.015625, 0.03125, 0.0625, \text{ and } 0.125$ ) QDs, CdS QDs, ZCS QDs, and ZnS QDs. (E) Electronic band structures of  $\text{Ni}_x\text{ZCS}$  ( $x = 0.015625, 0.03125, 0.0625, \text{ and } 0.125$ ) QDs, Pt-CdS QDs, CdS QDs, ZCS QDs, and ZnS QDs. (F) EIS Nyquist plots of  $\text{Ni}_x\text{ZCS}$  ( $x = 0.015625, 0.03125, 0.0625, \text{ and } 0.125$ ) QD, CdS QD, ZCS QD, and ZnS QD electrodes measured under the open-circuit potential of electrodes with visible-light irradiation. (G to J) Unfolded band structure of the supercell for bulk  $\text{Ni}_x\text{ZCS}$  ( $x = 0.015625, 0.03125, 0.0625, \text{ and } 0.125$ ) crystals, demonstrating the direct bandgap. The Fermi level moves to 0, as indicated by the dashed lines. Copyright 2020, American Association for the Advancement of Science.

mobility as well as conductivity of carriers, and adjusting the surface properties such as the interaction with reactants or intermediates.<sup>112-114</sup> In the photocatalytic water decomposition reaction, due to the lack of efficient catalytic active sites on the semiconductor surface, it is not only difficult to achieve effective adsorption and activation of reaction molecules, but also it seriously hinders the transfer of photo-generated charges to reaction molecules, which greatly limits the catalytic reaction efficiency. Therefore, defect construction on the surface of materials can not only provide active sites with coordination unsaturation for adsorption and activation of reactive small molecules, but also regulate the semiconductor energy band structure, accumulate photo-generated charges, and eliminate oxidation holes. At the same time, defects can be formed as part of the catalytic cycle and interact with the electrolyte.<sup>115-119</sup>

Wang *et al.* synthesized three dimensionally ordered macroporous (3DOM)  $\text{WO}_3$  with large holes by defect engineering

using  $\text{H}_2$  treatment.<sup>120</sup> The oxygen evolution rate of 3DOM  $\text{WO}_3$  (W270-400) was  $40.1 \mu\text{mol h}^{-1}$ , under visible light irradiation (300 W Xe lamp, cutoff at  $\lambda = 400 \text{ nm}$ ). It is much higher than that of untreated 3DOM  $\text{WO}_3$  and defect engineered  $\text{WO}_3$  nanoparticles, which was caused by good light collection and excellent charge transport, resulting from the formation of an amorphous layer with abundant surface oxygen vacancies on the catalyst surface. Ma *et al.* synthesized MOFs with controlled structural defects,  $\text{UiO-66-NH}_2\text{-X}$  ( $X$  is the molar equivalent of the regulator acetic acid relative to the linking agent during synthesis, Fig. 9).<sup>121</sup> With the increase of structural defects, the photocatalytic  $\text{H}_2$  production rate showed a volcanic trend. Among them,  $\text{Pt@UiO-66-NH}_2\text{-100}$  displayed the highest activity. Ultra-fine transient absorption spectrum analysis indicated that  $\text{UiO-66-NH}_2\text{-100}$  with moderate structural defects showed the fastest relaxation kinetics and the highest charge separation efficiency, while excessive defects delayed the

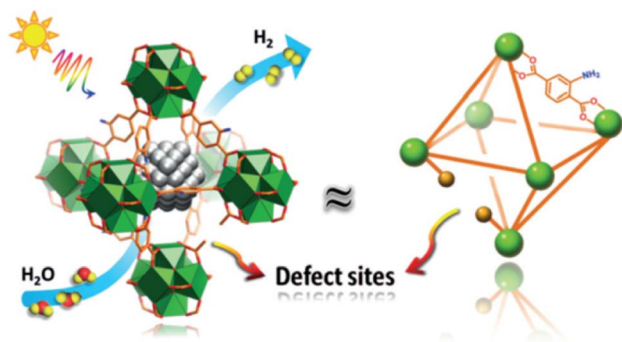


Fig. 9 Photocatalytic hydrogen production over Pt@UiO-66-NH<sub>2</sub>-X with structural defects. Copyright 2019, Wiley-VCH.

relaxation and reduced the charge separation efficiency. Therefore, the generation of moderate structural defects was crucial to promote the efficient separation of electrons–holes.

Vacancies are the main surface defects; however, the reports on vacancies in the crystal lattice are relatively few, especially for photocatalysis. Lin *et al.* prepared one-dimensional Cd<sub>1-x</sub>Zn<sub>x</sub>S@O-MoS<sub>2</sub>/NiO<sub>x</sub> hybrid nanostructures by adjusting zinc doping content, defect-rich O-MoS<sub>2</sub> layer and the growth of NiO<sub>x</sub> nanoparticles,<sup>122</sup> and optimized the energy band arrangement, morphology, surface active sites and interface charge separation to enhance the photocatalytic performance of hydrogen production. With the increase of Zn doping amount, HER activity of Cd<sub>1-x</sub>Zn<sub>x</sub>S ( $x = 0, 0.05, 0.10, \text{ and } 0.15$ ) increased, due to the increase of negative conduction potential. At the same time, after crossing Cd<sub>1-x</sub>Zn<sub>x</sub>S with defect-rich O-MoS<sub>2</sub> and NiO<sub>x</sub>, the H<sub>2</sub> formation rate of Cd<sub>1-x</sub>Zn<sub>x</sub>S @ O-MoS<sub>2</sub>/NiO<sub>x</sub> containing 15% Zn and 0.2 wt% O-MoS<sub>2</sub> (CZ0.15S-0.2 M-NiO<sub>x</sub>) reached up to 66.08 mmol h<sup>-1</sup> g<sup>-1</sup>, under visible light irradiation (a 300 W Xe lamp equipped with a 420 nm cutoff filter), because of the introduction of a large amount of active sites and the promotion of carrier separation, as shown in Fig. 10. Notably many different defects can exist simultaneously. A ZnIn<sub>2</sub>S<sub>4</sub> monolayer with double defects (Ag dopant and nanopore) was obtained by Pan *et al.* through cation exchange.<sup>123</sup> These two defects act as active sites for water oxidation (Ag dopant) and reduction (nanopore), which makes the ZnIn<sub>2</sub>S<sub>4</sub>

monolayer have optimized light absorption and carrier dynamics, resulting in stable photocatalytic overall water splitting performance without an auxiliary catalyst, under visible light irradiation ( $\lambda > 420 \text{ nm}$ ).

### S-scheme heterojunction

A photocatalyst can combine several semiconductors to form a heterojunction, which can broaden light collection characteristics and improve chemical stability, enhancing the separation of photo-generated carriers, reducing the recombination probability of photo-generated electrons and holes, and promoting photocatalytic water decomposition.<sup>124–131</sup> Straddling, staggered, and broken gaps have been intensively studied, and the Z-type heterojunction offers another solution for facilitating charge transfer in the junction. Notably, the Yu Group proposed a new type of heterojunction, S-scheme (or step-scheme) in 2019,<sup>132</sup> which could effectively realize the separation of electron–hole pairs with strong redox. Photocatalysts can be categorized into reduction photocatalysts (RPs) and oxidation photocatalysts (OPs), based on their CB and VB positions. RPs have high CB, while OPs have low VB. Therefore, photogenerated electrons are effective in RPs, while photogenerated holes are the contributive factors in OPs. An S-scheme heterojunction is composed of a RP and OP with staggered band structures, as indicated in Fig. 11. In the junction, the powerful photogenerated electrons and holes are reserved in the CB of the RP and VB of the OP, respectively, while the pointless photogenerated charge carriers are recombined, introducing a strong redox potential (Fig. 11). Although the structure is similar to type-II heterojunction, the charge-transfer route is totally different. If a charge-transfer route in a heterojunction goes with the S-scheme mechanism, the useless photogenerated charge carriers with weak redox ability are recombined with each other, while the useful photogenerated charge carriers with strong redox power are spatially separated and reserved, endowing it with strong redox ability and high charge-separation efficiency.<sup>57</sup> As a result, the S-scheme system possesses strong redox ability, which is endowed with a thermodynamic advantage in overall water splitting reaction. Since then, S-scheme heterojunctions have attracted more and more attention.



Fig. 10 Energy-band alignments of CZ<sub>0.15</sub>S, O-MoS<sub>2</sub>, and NiO<sub>x</sub> (left) and a schematic showing the photocatalytic hydrogen evolution over CZ<sub>0.15</sub>S@O-MoS<sub>2</sub>/NiO<sub>x</sub> (right image). Copyright 2019, Wiley-VCH.





Fig. 11 Band structure configuration and charge-transfer route of the S-scheme heterojunction. Copyright 2020, Elsevier Inc.

The investigation on the junction of ultrathin  $\text{WO}_3/\text{g-C}_3\text{N}_4$  nanosheets generates the first report on S-scheme heterojunctions, whose photocatalytic hydrogen production rate is  $\sim 1.7$  times higher than that of pure  $\text{g-C}_3\text{N}_4$ .<sup>132</sup> With the help of X-ray photoelectron spectroscopy, work function and electron paramagnetic resonance characterization, the enhanced photocatalytic performance was ascribed to the formation of the S-scheme heterojunction, which inhibited the recombination of useful electrons and holes, but expedited the recombination of relatively useless electrons and holes. Then many S-scheme heterojunctions are constructed, such as oxide/oxide (e.g.,  $\text{NiFe}_2\text{O}_4/\text{TiO}_2\text{-Ag}^+$ ),<sup>133</sup> oxide/sulfide (e.g.,  $\text{ZnWO}_4\text{-ZnIn}_2\text{S}_4$ ,  $\text{W}_{18}\text{O}_{49}/\text{Mn}_{0.2}\text{Cd}_{0.8}\text{S}$ ,  $\text{O-ZnIn}_2\text{S}_4/\text{TiO}_{2-x}$ ,  $\text{MoS}_2@\text{MoO}_3$ , and  $\text{Ti}_3\text{C}_2$  MXene@ $\text{TiO}_2/\text{CuInS}_2$ ),<sup>134-138</sup> MOF/oxide (e.g.,  $\text{Ni-MOF-74}/\text{BiVO}_4/\text{P}$  and  $\text{Ni-MOF-74}/\text{In}_2\text{O}_3$ )<sup>139,140</sup> and layered double hydroxides (e.g.,  $\text{CoS}_x/\text{NiAl-LDH}$  and  $\text{ZIF-67}@\text{NiAl-LDH}$ ).<sup>141,142</sup> For instance, the junction of  $\text{NiFe}_2\text{O}_4/\text{TiO}_2\text{-Ag}^+$  had enhanced light absorption with an absorption edge of 457 nm, and its maximum hydrogen production rate was  $27.4 \mu\text{mol g}^{-1} \text{h}^{-1}$ , under simulated sunlight illumination (HAL-320 Compact Xenon Light Source Solar Simulator,  $1000 \text{ W m}^{-2}$ ), owing to enhanced light absorption and mitigated  $e^-/h^+$  recombination, originating from the S-scheme heterojunction.<sup>133</sup> Zhang *et al.* reported *in situ* partial sulfidation of  $\text{MoO}_3$  powder to obtain a  $\text{MoS}_2@\text{MoO}_3$  S-scheme heterojunction photocatalyst.<sup>136</sup> The introduction of sulfur atoms could replace oxygen atoms to build a sulfur-rich surface and generate molybdenum sulfide. The hydrogen production rate was  $12\,416.8 \mu\text{mol h}^{-1} \text{g}^{-1}$ , illuminated with 300 W xenon light. Dai *et al.* successfully synthesized an S-scheme  $\text{ZnWO}_4\text{-ZnIn}_2\text{S}_4$  heterojunction with a two-dimensional coupling interface by a simple solvothermal method.<sup>134</sup> The  $\text{H}_2$  evolution activity of the  $\text{ZnWO}_4\text{-ZnIn}_2\text{S}_4$  composite with good stability reached  $4925.3 \mu\text{mol h}^{-1} \text{g}^{-1}$ , illuminated under 500 W xenon light. Chen *et al.* constructed S-doped  $\text{g-C}_3\text{N}_4$  (SCN) and N-doped  $\text{MoS}_2$  (NMS) ladder heterojunctions by one-step thermal polycondensation.<sup>143</sup> Within 4

hours of illumination under a 300 W xenon lamp, the hydrogen production rate of the material reached  $658.5 \mu\text{mol}^{-1} \text{h}^{-1}$ , which was about 23 times and 38 times higher than that of pure SCN ( $28.8 \mu\text{mol}^{-1} \text{h}^{-1}$ ) and NMS ( $17.4 \mu\text{mol}^{-1} \text{h}^{-1}$ ), respectively.

It is worth noting that the combination of junctions with other structural tuning approaches has attracted intensive studies. For example, Ruan *et al.* proposed a double S-scheme heterojunction, composed of  $\text{g-C}_3\text{N}_4$  nanosheets with hydrogen-doped rutile  $\text{TiO}_2$  nanorods and anatase  $\text{TiO}_2$  nanoparticles (Fig. 12).<sup>144</sup> The photocatalytic hydrogen evolution rate of the catalyst at 365 nm reached  $62.37 \text{ mmol}^{-1} \text{h}^{-1}$ , irradiated with a 300 W Xe lamp (200–1000 nm), and the apparent quantum efficiency was up to 45.9%. The remarkable improvement of photocatalytic performance was ascribed to the unique double S-shaped structure, which induced efficient charge separation and transfer. Another example is the combination of the Schottky junction and S-scheme heterojunction.  $\text{Ti}_3\text{C}_2$  MXene@ $\text{TiO}_2/\text{CuInS}_2$ , integrating Schottky/S-scheme heterojunction, was synthesized by a two-step hydrothermal process.<sup>138</sup> It exhibited synergistic effects for enhanced visible-light harvesting, improved charge transfer and increased active sites for photocatalytic  $\text{H}_2$  evolution. As a result, it delivered a  $\text{H}_2$  evolution rate of  $356.27 \mu\text{mol g}^{-1} \text{h}^{-1}$ , illuminated with a 300 W xenon lamp. Wang *et al.* constructed an  $\text{O-ZnIn}_2\text{S}_4/\text{TiO}_{2-x}$  photocatalyst by combining the S-scheme heterojunction with oxygen vacancies.<sup>136</sup> Owing to the synergistic effect of oxygen defects and the S-scheme heterojunction, a tailored energy band structure alignment with higher redox potentials and a larger Fermi level potential difference was achieved, resulting in more efficient S-scheme interface charge transfer and separation efficiency. As a result, the photocatalyst showed a  $\text{H}_2$  production rate of  $2584.9 \mu\text{mol g}^{-1} \text{h}^{-1}$  under AM1.5G illumination.

### Limitations of the structural tuning strategy

Currently SAs, defect control and S-scheme heterojunctions are the hot topics in related research areas including solar-driven hydrogen production. Great advancements have been achieved in the comprehension of the physicochemical properties of structurally tuned semiconductors or heterojunctions. Limitations, however, also exist for each tuning approach, and they are strongly time-dependent, because the shortcomings will be resolved or changed with deep investigations on the structure–activity relationship and the development of synthesis methods. These shortcomings are also important further research directions.

First complexity that all structural tuning approaches are facing is the potential structural reconstruction, which will induce defects such as strain and grain boundaries. It is well-known that strain and grain boundaries as well as reconstructed structures are all highly active centers. Therefore, the understanding on the mechanism for the enhancement of photocatalytic performance and the clarification of how SACs, defects or heterojunctions contribute in a catalyst are extremely crucial in future research. Another challenge is that three essential steps in photocatalytic processes are not isolated. For



Fig. 12 Schematic illustration of (a) the Z-scheme in natural photosynthesis, (b) Z-scheme in an artificial photosynthetic system, (c) S-scheme, and (d) twin S-scheme artificial photosynthetic system before and after contact, and under illumination. (e) Schematic diagram of electrostatic self-assembly of material. Copyright 2022, Wiley-VCH.

example, the enhancement of light absorption by vacancy control may induce the increase of charge recombination.<sup>145</sup> Nonetheless, the overall performance could still be improved if the extent of light absorption enhancement is much higher than that of charge recombination. This problem is general in all photocatalytic systems.

The metals in SACs are mostly rare and expensive; the production cost to the final form will be a challenge. When the SACs can be achieved in a cost-effective and mass producible way, it can be expected that they can play an important role in practical photocatalytic applications. Most recently, researchers have done some advancements in this topic, which shows high potential for resolving this issue. For instance, our recent work<sup>146</sup> demonstrated that Ir SACs could be easily synthesized on  $\text{Co}_3\text{O}_4$  nanowires using a simple dip coating and sol-flame annealing method. The total process lasts no more than 100 seconds without any instruments. This method could also be applicable to other metals (Fig. 13).

Moreover, the increase of the SAC content (>10 wt%) will undoubtedly enhance the photocatalytic activities, but the stability will be compromised, because the high surface energy of SACs will induce the rapid aggregation or the reaction with the surrounding reactant to form other undesired species.

SACs with single metals cannot provide the sites of multiple metal atoms or metal ensemble required by some catalytic reactions, which limits their applications. In contrast, classic metal nanoparticles can offer the sites of multiple metal atoms or metal ensemble, but they have low metal utilization efficiency compared with SACs. In this case, many research studies focus on the subnanometer nanoclusters rather than SACs.

A significant gap remains between the experimental and computational studies on SAC involved semiconductors or devices. Density functional theory (DFT) is generally an extremely helpful, powerful, and reliable tool to provide fundamental understanding of electrocatalysis. Unfortunately,

SACs are a real challenge for simulations, due to their elusive nature at the boundary between homogeneous and heterogeneous catalysis. When treated as homogeneous catalysts, SACs are individual atoms, whose chemical reactivity directly relates to their electron configuration and orbital diagram. In contrast, when treated as heterogeneous catalysts, SACs are supposed to stay put through the reaction's cycle, due to strong interaction with the underlying support, which in turn influences their charge and spin state, as well as their chemical reactivity. Authentically modeling SAC-support bonds is highly demanded for the accurate prediction of electronic and band structures. However, this modeling is also affected by the real operating conditions, which often differ from the low-temperature or high-vacuum conditions for DFT calculations. The complexity

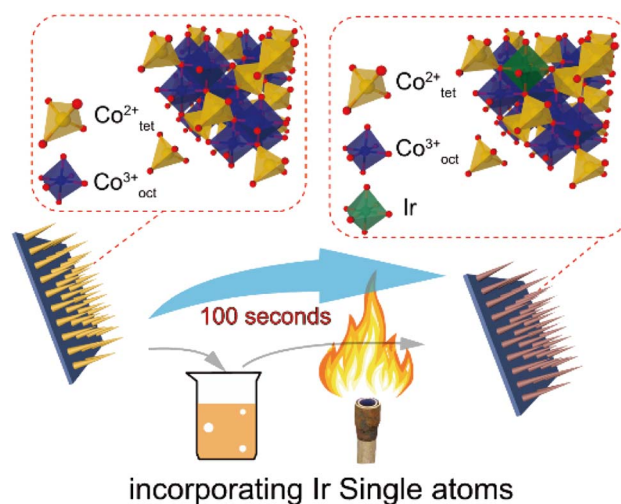


Fig. 13 Schematic illustration of synthesizing Ir SA incorporated  $\text{Co}_3\text{O}_4$  NWs. Copyright 2022, Wiley-VCH.

makes the DFT calculation highly correlated with the selected spin polarization and adopted DFT functional.

The limitation of computational studies on SACs is also applicable to defect control and heterojunctions because inaccurate structural modeling will provide meaningless or even false understanding on the structure–activity relationship. Defects such as vacancies that could be anywhere in the crystal lattice and in multiple forms make the DFT calculations extremely difficult, owing to the challenges in structural characterization on defects as well as the limitation in computer processors for processing an extremely large super cell. Existing characterization techniques can only provide indirect information about the atomic structure of defects at low resolutions, which hinders the establishment of the structure–activity relationship in photocatalytic reactions.

To boost the photocatalytic activities, bulk, surface, and interfacial defects should be finely controlled. Unfortunately, the existing synthetic methods are incapable of achieving defect control at such levels and can only tune one parameter of defects. Only when the defects are precisely controlled in photocatalytic materials can the rational defect design come into reality. More importantly, the defects such as vacancies could evolve during the photocatalytic process. For instance, bulk defects could evolve to surface defects and the latter could participate in the photocatalytic reactions. Therefore, how to measure the evolution of defects also remains a challenge. Present *in situ* and operando Raman, X-ray powder diffraction and infrared spectroscopy still have limitations at spatial, spectral, and temporal resolutions.

The S-scheme heterojunction is a milestone that researchers attempt to understand the charge transfer in the heterojunctions. It is a proposed new interpretation, but its scope is mainly limited to powder photocatalysts. It cannot be applied for PEC hydrogen production and solar cells with an external circuit until now. In addition, the S-scheme route is mainly used for the junction composed of two n-type semiconductors, which should have suitable band structures and obvious Fermi level differences.

## Conclusion and outlook

A promising photocatalytic hydrogen production system should have the following features: (1) high light absorption ability for visible and near infrared light. High light absorption coefficient is the same important as narrow bandgap. (2) Low recombination rate of photo-excited charge carriers. The charge pair transportation length should be higher than light absorption depth. (3) Fast surface redox reaction kinetics as well as decent stability. To achieve the goal, photocatalytic system designs especially on the composition and structure are highly demanded, and new composition and novel structures should be proposed. Given that it is a dramatic challenge for single semiconductors to possess all the features, a multi junction with engineered architectures is necessary to achieve high solar to hydrogen conversion efficiency. In addition, owing to the highly active nature of SACs, the application of SACs in photocatalysis will be more intensively studied. Herein a short

overview on the development of SAs, defect control and S-heterojunctions is given, and tries to inspire more novel ideas in the research on structural engineering approaches. In addition, from our point of view, further work will put more efforts in the following aspects. The clarification of the charge transfer mechanism, which is closely related to the catalytic performance. The careful and precise control of the band structure and electronic structures among the multiple junctions will be developed based on the clarification. The massive production of SAs with multiple elemental components and their applications for photocatalytic hydrogen production will be an important direction. And the synergetic effects between SAs with semiconductor substrates will be deeply studied using advanced experimental and theoretical methods, since the related topics are barely investigated. Active centers should be identified, which is crucial for interpreting the photocatalytic mechanism and developing more active and stable photocatalytic systems. Since the water splitting to produce hydrogen involves multiple electron transfer, the surface structure will definitely experience structural transformation. How to control the defects to make it both active and stable will be a hot topic. SACs with multiple elemental components should be developed to bridge SACs and subnanometer nanoclusters, and they could be ideal catalyst candidates because they can not only provide multiple metal atom sites, but also bear higher surface atom ratios than nanocluster counterparts. The structure of S-scheme photocatalysts needs to be rationally designed. The interface, junction morphology and band structures of each semiconductor should be precisely controlled. The interaction among various engineering strategies should be identified so as to effectively enhance the photocatalytic hydrogen evolution performance. To deepen the understanding on the structure–activity relationship and pinpoint the contribution of each structural tuning strategy, advanced *in situ* and *operando* observations at higher spatial, spectral, and temporal resolutions or multiple characterizations should be developed. Overall, the rational structural design strategy promotes quick advancements in photocatalytic water splitting and still needs further intensive studies.

## Author contributions

F. Xu, and B. Weng devised and outlined the draft of the manuscript.

## Conflicts of interest

There are no conflicts of interest.

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