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CdSe/ZIF-8-x: synthesis and photocatalytic CO₂ reduction performance†

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The photocatalytic reduction of CO₂ is an effective way to solve the greenhouse effect. Different kinds of materials, such as semiconductors, coordination compounds, and bioenzymes, have been widely investigated to increase the efficiency of the photocatalytic reduction of CO₂. However, a high selectivity and great stability are still challenges for material scientists. Here, we report for the first time visible light photocatalytic CO₂ reduction by a series of CdSe/ZIF-8 nanocomposites combining the excellent CO₂ adsorption capacity of ZIF-8 and the narrow energy gap of CdSe quantum dots (QDs). The composites show a higher catalytic performance than those of the pure components. Among CdSe/ZIF-8-x ($x = n_{\text{CdSe}}/n_{\text{ZIF-8}}$), the highest yield (42.317 $\mu\text{mol g}^{-1}$) for reducing CO₂ to CO in 12 h, was obtained using nanocomposites with a ratio of 0.42 ($n_{\text{CdSe}}/n_{\text{ZIF-8}}$) within the range of investigation.

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Nowadays, global energy shortage and environmental pollution are two major obstacles to the development of human society, and have attracted increasing concern. Using solar energy to convert CO₂ into valuable fuels or chemicals is extremely attractive due to its dual function of the reduction of the greenhouse effect and also as an alternative energy source to fossil fuels. Recently, different kinds of materials, such as semiconductor materials,^{1–3} metal complexes,^{4–6} and bioenzyme catalysts,⁷ have been explored for photocatalytic CO₂ reduction.

Metal-organic frameworks (MOFs) constructed from metal-containing clusters and organic building blocks are types of crystalline porous materials, and have been widely applied in many fields, such as gas storage,⁸ electrochemical energy storage (EES)^{9,10} and catalysis.¹¹ Recently, MOFs^{12–14} have been considered as potential new catalysts due to their excellent capability for CO₂ adsorption and capture.¹⁵ These porous materials provide a large number of catalytic active sites, and their porous structures are conducive to charge transfer.¹⁶ During the adsorption process, CO₂ coordinates with unsaturated metal sites and forms chemical bonds with MOFs.¹² Blom and co-workers demonstrated that CO₂ can interact with metal ions and form end-on adducts with one of the oxygen lone pair orbitals.¹⁷

ZIF-8, which is constructed from Zn²⁺ centres and imidazolate ligands, shows a high CO₂ adsorption capacity since the imidazolate ligand has a high adsorption capacity for CO₂ and also a strong complexation ability of CO₂.¹⁸ However, ZIF-8 has

a wide band gap (4.9 eV, ref. 19), which means that ZIF-8 is barely photoactive enough to catalyse CO₂ reduction. However, CdSe QDs can easily be excited to generate electron-hole pairs upon visible light irradiation due to their narrow band gap. Osterloh and co-workers reported CdSe QDs of several sizes applied to photocatalytic H₂ evolution and showed the quantitative relationship between the degree of quantum confinement and the photocatalytic H₂ evolution.²⁰

In this work, we synthesized a series of CdSe/ZIF-8-x composites, which combine the excellent CO₂ adsorption capacity of ZIF-8 with the narrow energy gap of CdSe QDs. X-ray diffraction (XRD) and energy dispersive spectroscopy (EDS) indicated the successful combination of CdSe QDs and ZIF-8. The CdSe/ZIF-8 composite exhibits an increased yield for reducing CO₂ to CO compared with pure CdSe QDs or ZIF-8. Under visible light irradiation for 12 h, the CO yield was 42.317 $\mu\text{mol g}^{-1}$, which is 6.13 and 10.84 times the yields catalysed by CdSe (6.901 $\mu\text{mol g}^{-1}$) and by ZIF-8 (3.905 $\mu\text{mol g}^{-1}$), respectively.

Reagents used in this work were analytically pure and used without further purification. Powder X-ray diffraction (PXRD) analysis was performed using a Rigaku Dmax-2000 diffractometer equipped with a Cu K α ($\lambda = 0.15406$ nm) radiation source. The morphology of the catalysts was observed by transmission electron microscopy (TEM, JEOL JEM-2100F) operated at 200 kV. Scanning electron microscopy (SEM) pictures were prepared using a Hitachi scanning electron microscope S-4800. Elemental mapping was carried out by energy dispersive X-ray spectroscopy (EDS) on the same instrument. Inductively coupled plasma spectrometry (ICP, Cary5000) was used for multi-elemental analyses. The CO₂ adsorption behaviours of the catalysts were studied with physical adsorption apparatus (ASAP

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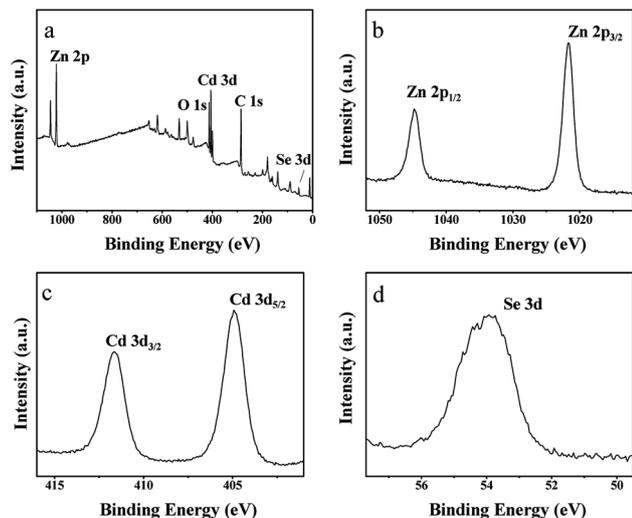


Fig. 3 (a) Survey spectrum of sample 2, (b) core level spectrum of Zn 2p, (c) core level spectrum of Se 3d, and (d) core level spectrum Cd 3d.

A typical EDS spectrum and elemental analysis of sample 2 are shown in Fig. S1 and Table S1,[†] respectively, confirming the presence of Cd, Se, Zn, C and O. The elemental ratio of $n_{\text{Cd}^{2+}}/n_{\text{Zn}^{2+}}$ calculated by the EDS is only 0.09, which is lower than that of the ICP result. This is probably due to the fact that the analysis of EDS comes from the surface elements and the lower elemental ratio indicates that CdSe is wrapped inside ZIF-8.

Fig. 3 shows the XPS survey spectrum and high-resolution spectra for Cd^{2+} 3d, Zn^{2+} 2p, and Se^{2-} 3d. As shown in Fig. 3(b), the $2p_{3/2}$ and $2p_{1/2}$ binding energies of Zn^{2+} are located at values of 1044.8 and 1021.7 eV, respectively. Fig. 3(c) shows the 3d peak of Se^{2-} at 54.1 eV. In addition, Fig. 3(d) shows that only two peaks appear, at binding energies of 411.7 and 404.9 eV, which are shifted towards the lower binding energy by about 0.3 eV of those of Cd^{2+} ($3d_{5/2}$) and Cd^{2+} ($3d_{3/2}$), from data reported in the literature.³⁰ The above results confirm the strong combination of CdSe and ZIF-8.

As shown in Fig. 4, among samples 1–3, sample 2 exhibits the highest CO_2 uptake at 298 K, which is about 13 times that of

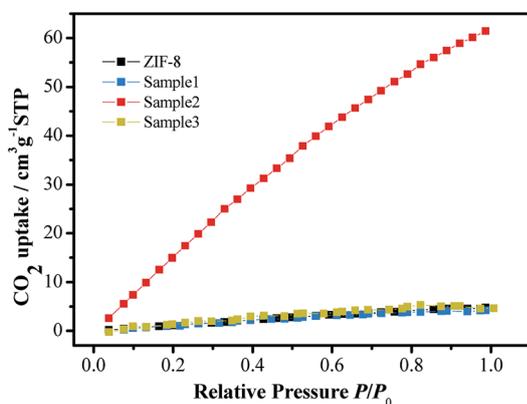


Fig. 4 CO_2 adsorption behaviour of CdSe/ZIF-8-x at 298 K.

ZIF-8. This result means that sample 2 can greatly absorb CO_2 before the reduction reaction so that it can accelerate the kinetic process of CO_2 reduction. Additionally, all the samples show a linear relationship between CO_2 uptake and relative pressure (0.1–1.0), indicating that the interaction between CO_2 and the samples is obviously physical.^{27,31} According to the DLS result, the greater adsorption performance of sample 2 could be due to the relatively uniform dispersion of CdSe in ZIF-8. While the aggregation of CdSe/ZIF-8-x results in a lower CO_2 uptake by the lower valid surface area and active sites from the unsaturated metal sites.¹² In addition, the morphology of the samples characterized by TEM, as shown in Fig. 2, indicates that the greater adsorption performance of sample 2 is due to the sufficient quantity of CdSe and the relatively uniform dispersion of CdSe in ZIF-8 and a lower aggregation of CdSe/ZIF-8-x.

Solid UV-visible diffuse reflectance spectroscopy (UV-vis DRS) was used to evaluate the band gap energy (E_g) of samples 1–3.³² UV-vis DRS of CdSe/ZIF-8 with different proportions were studied at room temperature. From Fig. 5(a), it can be seen that the absorption wavelength of ZIF-8 is about 302 nm, which is not in the visible light region. However, the visible light absorption ability of CdSe/ZIF-8 is obviously better than that of pure ZIF-8, and the spectral response range widened to 527–630 nm. In addition, we found that the E_g value of ZIF-8 is 4.88 eV, Fig. 5(b₁), which is too large to be used for visible light catalysis. However, the CdSe/ZIF-8 composites have much smaller E_g values than that of ZIF-8, at around 2.0 eV (Fig. 5(b₂–b₄)). This result supports the conclusion that the CdSe/ZIF-8 composites show better photocatalytic ability than ZIF-8.

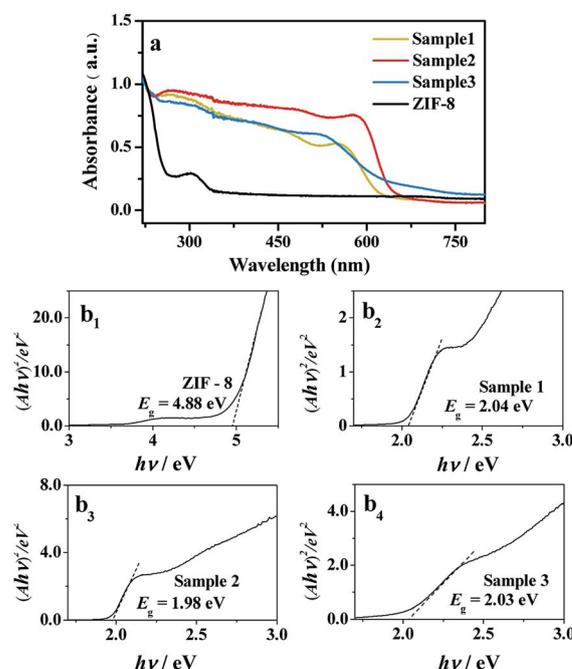


Fig. 5 (a) UV-vis DRS and (b₁–b₄) $(Ahn)^2$ vs. hn curves of CdSe/ZIF-8-x.



Table 2 Summary of CO₂ adsorption properties and photocatalytic activities of CdSe/ZIF-8-*x* (with CO₂ as the gas feedstock)

Sample	CO ₂ uptake (cm ³ g ⁻¹ STP)	CO production rate (μmol g ⁻¹ h ⁻¹)	CH ₄ production rate (μmol g ⁻¹ h ⁻¹)
ZIF-8	4.805	0.325	0.000
CdSe	—	0.575	0.000
Sample 1	4.219	1.521	0.092
Sample 2	61.435	3.526	0.102
Sample 3	4.655	2.038	0.056

To study the thermal stability of CdSe/ZIF-8-*x*, we conducted TGA. As shown in Fig. S3,[†] the initial decomposition temperatures of ZIF-8 and samples 1–3 are 118, 186, 422, and 158 °C, respectively, indicating that sample 2 has the highest thermal stability. Combined with the TEM results, the relatively uniform dispersion in sample 2 achieved the strongest combination force between CdSe and ZIF-8 among all the samples. Photocatalytic CO₂ reduction experiments were carried out under visible light irradiation, and the results are summarized in Table 2. Pure CdSe and ZIF-8 show low photocatalytic activities for CO₂ reduction, in which the CO production rate is below 1 μmol g⁻¹ h⁻¹. With an increase in the addition of CdSe, the rate of CO production first increases and then decreases. Among all the as-prepared samples, sample 2 shows the highest CO (3.526 μmol g⁻¹ h⁻¹) and CH₄ production rates (0.102 μmol g⁻¹ h⁻¹), in which the CO production rate was about 11 times that of ZIF-8 (0.325 μmol g⁻¹ h⁻¹) and about 6 times that of CdSe (0.575 μmol g⁻¹ h⁻¹). However, when using N₂ to replace CO₂ to start the reaction under identical conditions, a little CO and CH₄ were detected. The control experiment indicated that the CO product comes from CO₂ gas (Table S2[†]).

In a photocatalytic process, CO₂ adsorption is the rate-limiting step,³³ which is attributed to the fact that the CO₂ conversion efficiency of the photocatalyst significantly relies on the amount of CO₂ molecules adsorbed.¹⁶ Combined with the DLS result, the higher CO₂ uptake of sample 2 is the main reason for the high photocatalytic activities for CO₂ reduction. Theoretically, increasing the yield of CH₄ is more difficult than that of CO, because reducing CO₂ to CO consumes two electrons while eight electrons are needed in the CH₄ transformation.

Given the experimental results that CdSe/ZIF-8-*x* showed a higher CH₄ production rate than pure CdSe and ZIF-8, we considered that ZIF-8 pores could play the role of a “nano-reactor” to enclose CO₂ and CO, so as to finish the lengthy transformation process and improve the yield of CH₄.³⁴ In addition, the above inference could be supported by the CO₂ adsorption capacities data (Table 2).

Based on all the above results, the probable mechanism for CdSe/ZIF-8-*x* in the photocatalytic process is proposed as follows (Fig. 6). As shown in Fig. 6, CdSe QDs play the core role in the photocatalytic process, as they were excited to generate electron–hole pairs upon visible light irradiation due to their narrow band gap. Furthermore, the addition of CdSe improved the conductivity of CdSe/ZIF-8-*x*, which attributed to the charge transfer since a good conductivity leads to only a small charge transfer resistance.^{35,36} ZIF-8 plays a key role as the “electrons transporter” and also as the “nanoreactor”, which means that photogenerated electrons could be transferred quickly from CdSe to ZIF-8. Then, the molecular [Ru(bpy)₃]²⁺ photosensitizer can effectively receive the photoinduced electrons to reduce the CO₂ molecule absorbed by the ZIF-8 pores to yield CO. On the other hand, the photogenerated holes are quenched by TEOA acting as a sacrificial electron donor.

Conclusions

In this study, a series of CdSe/ZIF-8-*x* nanocomposites was synthesized. They have a higher photocatalytic activity than pure CdSe and ZIF-8. PXRD, TEM and EDS results indicate that the CdSe QDs were successfully combined with ZIF-8. Moreover, sample 2, in which $n_{\text{Cd}^{2+}}/n_{\text{Zn}^{2+}}$ is equal to 0.4, shows a higher thermal stability and increased yield for reducing CO₂ to CO (3.526 μmol g⁻¹ h⁻¹) and CH₄ (0.102 μmol g⁻¹ h⁻¹). The CO yield is about 11 times that of ZIF-8 (0.325 μmol g⁻¹ h⁻¹).

Conflicts of interest

There are no conflicts to declare.

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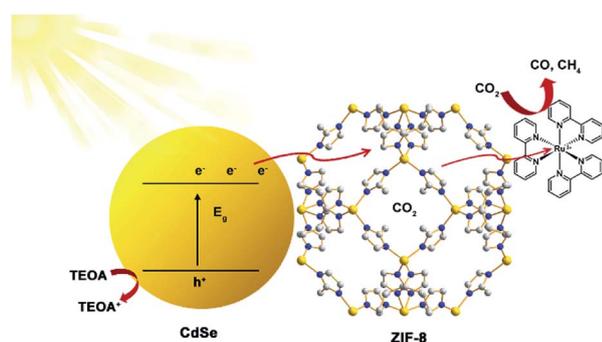


Fig. 6 Schematic illustration of the proposed mechanism of photocatalytic CO₂ reduction over CdSe/ZIF-8-*x*.



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