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Enhancing interfacial charge transfer in mesoporous MoS₂/CdS nanojunction architectures for highly efficient visible-light photocatalytic water splitting†

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Modulating the heterojunction interfaces at the nanoscale is paramount to developing effective photocatalysts. Nanodimensional heterostructures, essentially, expose a large fraction of active edge sites, which, along with excellent electronic communication, can result in high solar energy to chemical fuel conversion efficiencies. In this work, mesoporous heterojunction architectures made of ultrasmall MoS₂ nanolayers (ca. 10–15 nm in lateral size) and sub 5 nm sized CdS nanocrystals have been prepared through a polymer-templated oxidative aggregation of CdS nanocrystals followed by a wet-chemical deposition of exfoliated MoS₂ nanosheets on the CdS surface. Thorough chemical, morphological and structural investigations by electron microprobe analysis, X-ray diffraction, electron microscopy, X-ray photoelectron spectroscopy and N₂ physisorption prove that the MoS₂-modified CdS nanocatalysts are composed of a porous network of connected cubic CdS nanocrystals and 2H-phase MoS₂ nanosheets and possess a high internal BET surface area (ca. 159–225 m² g⁻¹) and uniform pores (ca. 6–9 nm in diameter). Photocatalytic studies coupled with UV–vis/NIR, photoluminescence and electrochemical impedance spectroscopy measurements indicate that the nanoscale MoS₂/CdS junctions provide more efficient electronic connectivity and charge carrier dissociation across the catalyst interfaces, resulting in a remarkable enhancement in the photocatalytic H₂ production activity. The optimized MoS₂/CdS catalyst at 20 wt% MoS₂ content achieves a H₂ production rate up to ~0.4 mmol h⁻¹ (or ~19 mmol h⁻¹ g⁻¹ mass activity) with remarkable stability under visible light irradiation, corresponding to an overall 6.7x enhancement of H₂ generation efficiency relative to the unmodified CdS. We also obtained an apparent quantum yield (AQY) of 51.2% for the hydrogen generation reaction using monochromatic light of 420 nm.

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1. Introduction

Over the last few years, various renewable technologies have been used to effectively produce chemical fuels, such as hydrogen which is a promising alternative target for clean chemical energy source.^{1,2} Among them, solar hydrogen production by water-splitting photocatalysis has emerged as a particularly propitious approach because of the low cost, easy maintenance and environmental effectiveness.³ Thus far, a great variety of semiconducting materials, including metal oxides⁴ and sulfides,^{5,6} have been applied for catalytic hydrogen fuel gene-

ration. These materials satisfy the most important criteria of effective photocatalysts, such as substantial photon absorption (bandgap energy ~2–2.8 eV) and favourable reduction and oxidation band positions.^{7–9} Metal sulfides, especially CdS, have been extensively studied due to their narrow bandgap (ca. 2.3–2.6 eV) and high negative conduction band (CB) level relative to the thermodynamic H⁺ reduction potential (–0.41 V, pH 7).^{10–12} However, the practical use of these materials is still plagued by photochemical corrosion and low photon-to-hydrogen conversion efficiency. To improve the photocatalytic activity of CdS, various strategies have been developed, including heterojunctions, core-shell structures, doping with metal or non-metal elements and loading of co-catalysts. For example, numerous reports have shown that electron-accepting co-catalysts, such as noble metals Pd and Pt on a CdS support, can mitigate charge recombination and thus accelerate the surface reaction for hydrogen evolution.^{13,14} Moreover, considering the high cost and limited availability of noble metals, in-

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expensive Earth-abundant metal-based co-catalysts have been recently examined.^{15,16}

After the revolution of graphene, transition metal dichalcogenides (TMDs), with the formula MX_2 where M is Mo, W, etc. and X is S, Se, and Te, have also received attention as ideal co-catalysts for the photocatalytic H_2 evolution reaction.¹⁷ Among TMDs, 2D molybdenum disulfide (MoS_2), consisting of stacked S–Mo–S layers weakly bonded by van der Waals forces, has attracted significant attention owing to the low cost and excellent optical, physical and electronic properties.¹⁸ In the bulk or multilayer form, MoS_2 has an indirect energy gap of about 1.3 eV, which transitions to a direct gap of 1.9 eV for a single monolayer. Besides the thermodynamically stable hexagonal 2H phase (semiconducting phase), MoS_2 also adopts two metastable phases, that is, the rhombohedral 3R (semiconducting phase) and trigonal 1T (metallic phase) structure.¹⁹ Among these polytypes, 2H- MoS_2 although exhibits lower mobility for transporting charge carriers than the metallic 1T- MoS_2 , shows better stability and interesting semiconducting properties that are useful for applications in light energy conversion and optoelectronics.²⁰ Also, the 2H phase of MoS_2 can exhibit better photocatalytic activities than the semiconducting 3R counterpart due to the unexfoliated structure of 3R- MoS_2 . In particular, the three-layered (3R) MoS_2 form adopts a relatively more stable stacking structure between the S–Mo–S layers, which can result in lower accessibility of the reactants to the inter-layer active sites.²¹ For this purpose, considering the semiconducting properties of MoS_2 , there have been extensive studies on various heterostructures using 2H- MoS_2 as the co-catalyst for improved H_2 production yield.^{22,23} For instance, Zhang *et al.* demonstrated the selective growth of single-layer MoS_2 nanoflakes, with a 4–10 nm lateral size, on the Cd-rich (0001) surface of wurtzite CdS particles.²⁴ This catalyst was reported to undergo photocatalytic reduction of water in the presence of lactic acid as the hole scavenger, yielding a H_2 evolution rate of $1.47 \text{ mmol h}^{-1} \text{ g}^{-1}$. Song's group successfully prepared a 2D/2D heterojunction by assembling ultrathin MoS_2 sheets and exfoliated CdS nanolayers through an adsorption–calcination process.²⁵ They found that the optimal photocatalyst presented a maximum photocatalytic H_2 production activity of $18.43 \text{ mmol h}^{-1} \text{ g}^{-1}$. More recently, Fujishima and coworkers coupled 2D MoS_2 thin sheets as H_2 evolution catalysts with CdS microspheres by a template-free solvothermal method.²⁶ The nanostructured core–shell CdS@ MoS_2 particles showed high carrier transport efficiency and a remarkable H_2 production activity of $4.16 \text{ mmol h}^{-1} \text{ g}^{-1}$ under natural solar light.

In the previous work, we have described the synthesis of mesoporous networks of small-sized CdS nanocrystals (NCs) through a polymer-assisted self-assembly method.²⁷ These materials compared to the isolated nanoparticles and conventional porous solids have shown adequate photocatalytic properties, which resulted from the strong electronic communication between the constituent nanoparticles and the large exposed surface area of the assembled structure. Such CdS NC-linked structures combine a number of interesting features, such as a 3D open-pore structure, high internal surface area,

visible light absorption and excellent structural stability.²⁸ In this study, we report the synthesis of new high surface-area mesoporous frameworks consisting of CdS NCs (*ca.* 4–5 nm in diameter) and exfoliated 2H- MoS_2 nanosheets (NSs) with an ultrasmall lateral size (*ca.* 10–15 nm) and evaluate their performance as catalysts for water splitting and H_2 evolution under visible light. The 2D/3D MoS_2 /CdS nano-heterostructures with various loadings of MoS_2 NSs, namely from 5 to 25 wt%, were synthesized through a facile deposition method. By using a combination of spectroscopic and electrochemical techniques, we provide an in-depth understanding of the structural, electronic and photochemical characteristics of the obtained catalysts. We demonstrate that the nanoscale dimension of the MoS_2 /CdS contacts minimizes electron–hole recombination and increases the electron density and mobility at the MoS_2 /CdS junction, yielding accelerated water reduction kinetics. Owing to these characteristics, the optimized catalyst with 20 wt% MoS_2 NSs exhibits a significant enhancement of the photocatalytic activity, yielding an apparent quantum yield (AQY) of 51.2% for H_2 evolution at 420 nm, a record efficiency for MoS_2 -modified CdS photocatalysts, to the best of our knowledge.

2. Experimental section

2.1 Synthesis of catalysts

2.1.1 CdS NC synthesis. CdS NCs of 4–5 nm size were synthesized according to ref. 29. In a typical procedure, CdCl_2 (7 mmol) was dissolved in deionized (DI) water (50 mL), to which 3-mercaptopropionic acid (3-MPA, 15 mmol) was added and the mixture was left under continuous stirring at room temperature (RT) for 15 min. The pH of the solution was then increased to 9–10 using concentrated NH_4OH . Next, a Na_2S aqueous solution (7 mmol in 20 mL) was gradually added and the resulting yellow colloidal sol was kept under stirring at RT for an additional 1 h. The MPA-stabilized CdS NCs were collected by centrifugation with the addition of 2-propanol (60 mL), and dried at 40 °C overnight.

2.1.2 Mesoporous CdS NCAs. The procedure to prepare mesoporous CdS NC assemblies (NCAs) was based on our previously reported work.²⁷ Briefly, 2.0 mmol MPA-stabilized CdS NCs were first suspended in DI water (2.5 mL) under stirring at RT. Then, few drops of NH_4OH were added until a clear colloid solution was observed. Next, this colloid sol was dropped into an aqueous solution containing Pluronic F-127 (~10% w/v, 2.5 mL) and the resulting mixture was kept under stirring at RT for 1 h. Finally, H_2O_2 solution (3 wt%, 1 mL) was added dropwise under continuous stirring until gelation was observed. After being stirred for at least 30 min, the gel suspension was transferred into a 50 mL glass beaker and heated in an oven for 3 days at 40 °C to slowly evaporate the solvent under static conditions. The final mesoporous product was collected after several washing cycles (one cycle with 20 ml of ethanol and three cycles with 20 ml of DI water), vacuum filtration and drying at 40 °C overnight.



2.1.3 MoS₂ NS synthesis. For the synthesis of the 2H-MoS₂ NSs, monovacant Keggin-type phosphomolybdate (Na₇[PMo₁₁O₃₉]) was synthesized following a previously reported procedure.³⁰ Next, Na₇[PMo₁₁O₃₉] (0.0175 mmol) was dissolved in DI water (10 mL) at RT. To this solution, thioacetamide (1 mmol) was added and the resulting mixture was kept under stirring at RT for 2 h. The final blue solution was sealed in a 50 mL Teflon-lined autoclave and heated in an oven at 200 °C for 12 h. The black product was isolated by centrifugation, washed with water and dried at 60 °C for 24 h.

For comparison purposes, MoS₂ microflakes were also synthesized following a similar procedure, but using (NH₄)₆Mo₇O₂₄·4H₂O (0.0275 mmol) as the Mo source.

2.1.4 Mesoporous MoS₂/CdS NCAs. Porous structures of MoS₂ NSs and CdS NCs with different MoS₂ contents, namely, 5, 10, 15, 20 and 25 wt%, were prepared by a facile deposition method. For a typical synthesis of a 20 wt% MoS₂-modified CdS sample, the as-prepared MoS₂ NSs (20 mg) were added to a H₂O : isopropanol (IPA) solution (2 : 1 v/v, 30 mL) in a 50 mL beaker. The suspension was ultrasonicated for 2 h until a homogeneous black solution of exfoliated MoS₂ NSs was formed. In a separate vial, mesoporous CdS NCAs (80 mg) were dispersed in a H₂O : IPA solution (2 : 1 v/v, 5 mL) under vigorous stirring at RT. Finally, the suspension of exfoliated MoS₂ NSs was added gradually to the CdS NCA dispersion and the resulting green mixture was left under stirring and low heat (50–60 °C) for slow evaporation of the solvents. The respective 5, 10, 15 and 25 wt% MoS₂-modified CdS samples were synthesized following a similar procedure using the stoichiometric amount of MoS₂ NSs for each case.

A bulk-like MoS₂-modified CdS catalyst (designated as 20-MoS₂/CdS_b) was also prepared by depositing 20 wt% of MoS₂ microflakes on the surface of mesoporous CdS.

2.2 Photocatalytic water splitting study

The photocatalytic H₂ evolution experiments were performed in an air-tight Pyrex glass cell with 20 mg of catalyst suspended in 20 mL of aqueous solution containing 0.25 M Na₂SO₃ and 0.35 M Na₂S (except otherwise noted). Before irradiation, the suspension was bubbled with argon for 40 min. The reactor was illuminated using a 300 W Xe lamp (Variat Cermax) with a UV-cutoff filter ($\lambda \geq 420$ nm) and the reaction temperature was thermostated to 20 ± 2 °C by a water-cooling system. During the experiment, 100 μ L of gas was collected from the reactor's headspace and the generated H₂ was measured using a gas chromatograph (Shimadzu GC-2014, TCD, with Ar as the carrier gas).

The apparent quantum yield (AQY) was estimated by measuring the amount of hydrogen evolved at $\lambda = 420 \pm 10$ nm irradiation wavelength, according to the following equation:

$$\text{AQY} = \frac{\text{(amount of evolved H}_2 \text{ molecules} \times 2)}{\text{(total incident photons)}}$$

The intensity of light was measured using a StarLite power meter equipped with a FL400A-BB-50 thermal sensor (Ophir Optonics Ltd).

2.3 Characterization methods

X-Ray diffraction was performed on a Panalytical X'pert Pro MPD instrument with Cu-K α radiation ($\lambda = 1.5418$ Å), operated at 45 kV and 40 mA. Energy-dispersive X-ray spectroscopy was performed on a JEOL JSM 6390 LV scanning electron microscope (SEM) equipped with an Oxford INCA PentaFETx3 detector (Oxfordshire, UK). For EDS analysis, because of the overlap of Mo-L α and S-K α lines, the samples were annealed at 800 °C in air for 3 h in order to detect the Mo : Cd atomic ratios, and the measurements were performed on at least five different regions of each sample using 20 kV acceleration voltage and 100 s accumulation time. Transmission electron microscopy measurements were performed with a JEOL JEM-2100 instrument (LaB₆ filament) operating at 200 kV. The samples were prepared by dispersing fine powders in ethanol using sonication and then drop-casting on a holey carbon-coated Cu grid. X-ray photoelectron spectroscopy (XPS) was conducted on a SPECS spectrometer using a Phoibos 100 1D-DLD electron analyzer and Al K α radiation as the energy source (1486.6 eV). Binding energy values were corrected with reference to the C 1s (284.8 eV) signal of adventitious carbon. UV-vis/near IR diffusion reflectance spectra were recorded with a Shimadzu UV-2600 spectrophotometer, using BaSO₄ powder as a 100% reflectance reference. The diffuse reflectance data were transformed with the Kubelka–Munk function [$\alpha/S = (1 - R)^2/(2R)$], where R is the measured reflectance and α and S are the absorption and scattering coefficients, respectively] into absorption spectra. N₂ adsorption–desorption isotherms were obtained at –196 °C with a Quantachrome NOVA 3200e system. Prior to analysis the samples were heated at 80 °C under low pressure (<10^{–5} Torr) to remove residual solvents and water. The surface areas were determined from the adsorption data (P/P_0 range 0.04–0.22) using the Brunauer–Emmett–Teller (BET) method, and the total pore volumes were calculated from the adsorbed N₂ amount at $P/P_0 = 0.98$. The pore size distribution plots were calculated from the adsorption data using the non-local density functional theory (NLDFT) fitting model. Room temperature photoluminescence and decay curve analysis were performed with an Edinburgh FS5 spectrofluorometer.

2.4 Electrochemical measurements

Mott–Schottky plots, Nyquist plots and current–voltage (J – V) curves were measured in a 0.5 M Na₂SO₄ electrolyte (pH = 7) using a single-channel potentiostat–galvanostat (Princeton Applied Research VersaSTAT 4) equipped with a three-electrode cell, consisting of a sample-coated FTO working electrode, an Ag/AgCl (saturated KCl) reference electrode, and a Pt-wire counter electrode. The working electrodes were prepared as follows: 10 mg of each sample were dispersed in a 2-propanol/DI water solution (1 : 2 v/v, 1 mL). Then, 20 μ L Nafion solution (5 wt%) was added and the mixture was ultrasonicated for 15 min and left under stirring for 24 h. When a uniform suspension was formed, 100 μ L of the solution was dropped on a fluorinated tin oxide (FTO, 10 Ω sq^{–1}) substrate (effective



surface 1 cm^2) and the film was dried under heat at $40\text{--}50 \text{ }^\circ\text{C}$ for 1 h.

The space-charge capacitance (C_{sc}) of the semiconductor/electrolyte interface was measured at 1 kHz, with 10 mV AC voltage amplitude and the obtained flat-band potentials were converted to the reversible hydrogen electrode (RHE) scale using the equation: $E_{\text{RHE}} = E_{\text{Ag/AgCl}} + 0.197 \text{ V}$, where E_{RHE} is the potential in the RHE scale and $E_{\text{Ag/AgCl}}$ is the measured potential in the Ag/AgCl scale.

The donor density (N_{D}) of the as-prepared materials was calculated according to the Mott-Schottky equation: $N_{\text{D}} = C_{\text{sc}}^2 \cdot 2 \cdot (E - E_{\text{FB}}) / \epsilon \cdot \epsilon_0 \cdot e_0$, where, C_{sc} is the space charge capacitance, E_{FB} is the flat band potential, E is the applied potential, N_{D} is the donor density of the electrode material, ϵ is the dielectric constant (8.9 for CdS), ϵ_0 is the vacuum permittivity ($8.8542 \times 10^{-14} \text{ F cm}^{-1}$), e_0 is the elementary charge ($1.602 \times 10^{-19} \text{ C}$) and the term $(E - E_{\text{FB}}) \cdot C_{\text{sc}}^2$ is the reciprocal of the slope of the Mott-Schottky plot.

For Nyquist plots, the current output was recorded between 1 Hz and 1 MHz with a 10 mV AC perturbation, under open-circuit potential. The EIS data were fitted to an equivalent circuit model using ZView Software.

3. Results and discussion

3.1 Materials synthesis and characterization

Porous networks of connected CdS NCs and 2H-MoS₂ NSs were produced using a polymer-templated oxidative aggregating method followed by a wet-chemical deposition process. Initially, 2D nanolayers of 2H-MoS₂ were prepared through a hydrothermal sulfurization reaction, using Na₇[PMo₁₁O₃₉] and thioacetamide (CH₃CSNH₂) as the Mo and S source, respectively.³¹ After annealing to a certain temperature, thioacetamide begins to decompose and evolve H₂S that is capable of transforming MoO_xⁿ⁻ species into MoS₂. The X-ray diffraction (XRD) pattern of as-made MoS₂ NSs, in Fig. S1,† indicates the successful chemical transformation of phosphomolybdate clusters into MoS₂ under our liquid-phase conditions; all the diffraction peaks can be assigned to the hexagonal (2H) structure of MoS₂. Besides, the broadened XRD diffractions point to the nanoscale dimension of the MoS₂ layers. Nanoporous networks of CdS NC assemblies (NCAs) were prepared based on a

method described in a recent study, which involves the oxidative coupling of thiol-capped CdS nanoparticles with the assistance of a tri-block copolymer (Pluronic F-127) surfactant. MoS₂ modification of the mesoporous CdS structure was realized by depositing the liquid exfoliated MoS₂ NSs onto the surface of CdS NCAs. Following this process, several *n*-MoS₂/CdS NCAs with different compositions of grafted MoS₂, namely *n* = 5, 10, 15, 20 and 25 wt%, were prepared. The Mo/Cd atomic ratios of the as-prepared materials were determined by elemental analysis with energy dispersive X-ray spectroscopy (EDS). All the EDS spectra affirm that the Mo content in MoS₂/CdS NCAs is very close to the nominal composition of the samples (within an 8% deviation), suggesting the complete binding of MoS₂ to the CdS surface (Fig. S2,† and Table 1).

The crystallinity and surface chemistry of the as-synthesized materials were examined by X-ray diffraction (XRD) and X-ray photoelectron spectroscopy (XPS). As shown in Fig. 1a, the XRD patterns of CdS and MoS₂/CdS NCAs display three Bragg diffraction peaks at 2θ scattering angles of $20^\circ\text{--}60^\circ$, which can be indexed to the (111), (220) and (311) diffractions of the zinc blende phase of CdS (JCPDS no. 42-1411). Besides, the peak widening of the XRD diffractions implies the formation of CdS particles with a very small grain size; indeed, peak-width analysis using the Scherrer equation gave an average CdS domain size of $\sim 2.3 \text{ nm}$. Note that, after MoS₂ deposition, CdS still maintains its crystallinity, indicating that MoS₂ nanolayers are mainly studded on the CdS surface. In XRD patterns, however, no additional peak is identified after MoS₂ deposition, even in the XRD plot of the composite sample with 25 wt% MoS₂, which is due to the good dispersion and nanoscale size of the MoS₂ layers.

Typical XPS spectra of the best catalyst (20-MoS₂/CdS) are shown in Fig. 1b–d. The XPS Cd 3d spectrum, in Fig. 1b, presents a doublet peak at 405.3 and $412.2 \pm 0.2 \text{ eV}$ binding energies, corresponding to the Cd 3d_{5/2} and Cd 3d_{3/2} core level lines of Cd²⁺ in CdS, respectively.³² The XPS spectrum of the Mo 3d scan confirmed the presence of Mo⁴⁺ species in MoS₂, showing two peaks at $229.1 \pm 0.2 \text{ eV}$ (3d_{5/2}) and $232.4 \pm 0.2 \text{ eV}$ (3d_{3/2}) binding energies with a spin-orbit splitting of 3.3 eV (Fig. 1c), in line with other reports.³³ Besides the prominent Mo⁴⁺ 3d signal, there exists a weak doublet peak at a relatively higher binding energy ($231 \pm 0.2 \text{ eV}$ and $234.4 \pm 0.2 \text{ eV}$), which

Table 1 Chemical composition and textural and optical properties of the as-prepared CdS and MoS₂/CdS NCAs

Catalyst	MoS ₂ content ^a (wt%)	BET surface area (m ² g ⁻¹)	Pore volume (cm ³ g ⁻¹)	Pore size (nm)	Energy gap (eV)
CdS NCAs	—	225	0.34	9.1	2.71
5-MoS ₂ /CdS	5.4	196	0.26	8.1	2.71
10-MoS ₂ /CdS	10.8	185	0.23	6.6	2.71
15-MoS ₂ /CdS	15.9	176	0.23	6.6	2.68
20-MoS ₂ /CdS	21.7	166	0.22	6.6	2.68
25-MoS ₂ /CdS	26.2	159	0.20	6.1	2.63
20-MoS ₂ /CdS _b	21.5	169	0.23	7.6	2.67

^a Based on EDS analysis.



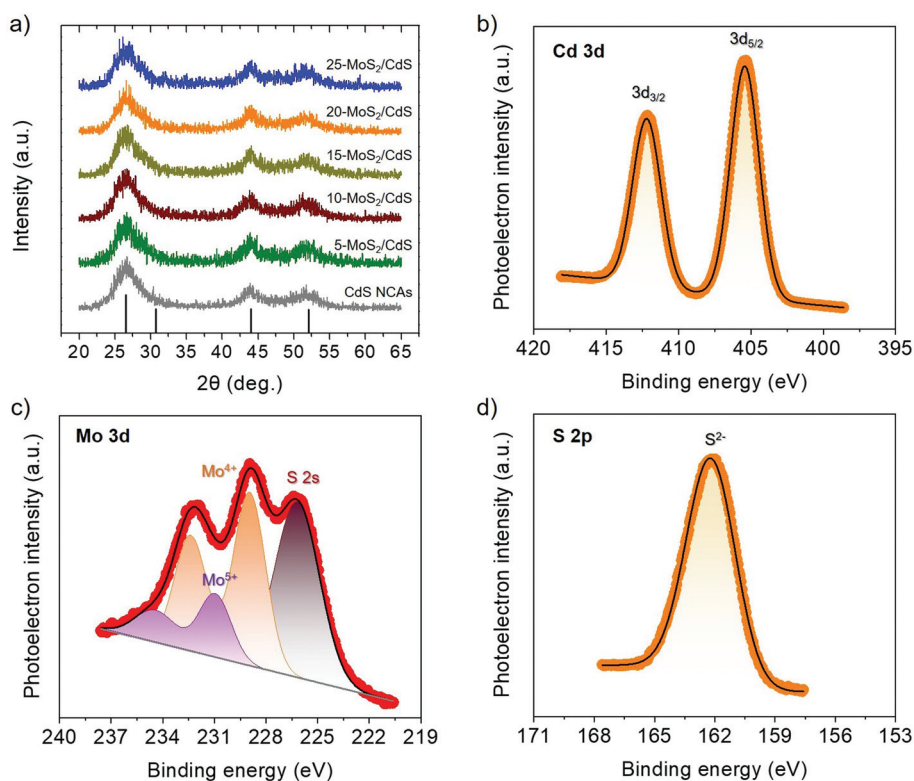


Fig. 1 (a) XRD plots for the mesoporous CdS and MoS₂/CdS NCAs. The standard diffraction data of the zinc blende CdS according to the JCPDS card no. 42-1411 are also shown (black bars). (b–d) Typical XPS core-level spectra of the Cd 3d (b), Mo 3d (c) and S 2p (d) region for the 20-MoS₂/CdS catalyst.

can be assigned to unsaturated Mo⁴⁺ atoms and/or higher oxidation state of Mo⁵⁺ species due to mild surface oxidation after the synthesis process.³⁴ As for the singlet peak observed at 226.6 ± 0.2 eV, this is associated with the S 2s signal of sulfides (S²⁻).³⁵ Furthermore, the XPS S 2p spectrum (Fig. 1d) exhibited a prominent peak at 162.2 ± 0.3 eV that is characteristic of the S²⁻ ions in metal sulfides.³⁶ Quantitative analysis of the XPS spectra also indicated a Mo-to-Cd atomic ratio close to 20.4 : 79.6 that corresponds to a ~ 22.1 wt% MoS₂ content, which is comparable to the composition obtained by EDS.

The morphology of the title materials was recorded *via* transmission electron microscopy (TEM), and the representa-

tive TEM images for the 20-MoS₂/CdS sample are shown in Fig. 2. The images reveal that the framework of 20-MoS₂/CdS is composed of approximately 4–5 nm sized CdS interconnected nanoparticles (Fig. 2a). In addition, from the high-resolution TEM (HRTEM) observation in Fig. 2b, some layered particles with a lateral dimension of around 10–15 nm can be discerned on the surface of CdS. A careful inspection reveals a layer-to-layer distance of ~ 6.5 Å that reasonably correlates to the (002) planes of MoS₂ NSs. This suggests a small enlargement of the interlayer distance of MoS₂ NSs with respect to the bulk MoS₂ (6.2 Å), which is due primarily to the small lateral size of the MoS₂ layers. A similar effect was previously observed for other MoS₂ nanostructures.³⁷ Taken together with XPS characterization of the surface structure, these results confirm that small-size MoS₂ NSs are anchored on the surface of linked CdS nanoparticles and, importantly, with a close contact that is advantageous for interfacial charge transfer and separation.

The N₂ physisorption method was applied to determine the porosity of the as-synthesized materials. The nitrogen adsorption and desorption isotherms and associated pore size distribution of the mesoporous 20-MoS₂/CdS NCAs are compared as shown in Fig. 3a to those of unmodified CdS NCAs. The respective plots for the other MoS₂-modified catalysts (*i.e.*, with 5, 10, 15 and 25 wt% MoS₂ content) are depicted in Fig. S3.† All samples display characteristic type-IV isotherms with a H₂-type hysteresis loop (according to the International

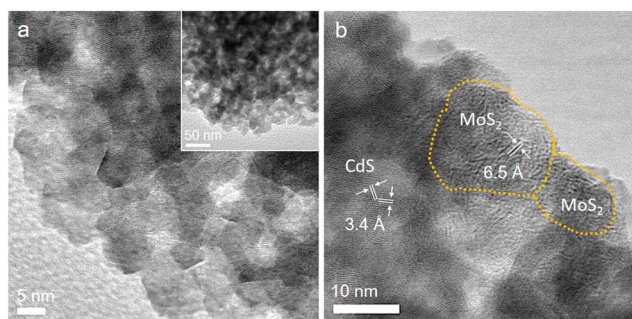


Fig. 2 Representative (a) TEM and (b) HRTEM images of the 20-MoS₂/CdS NCAs catalyst.



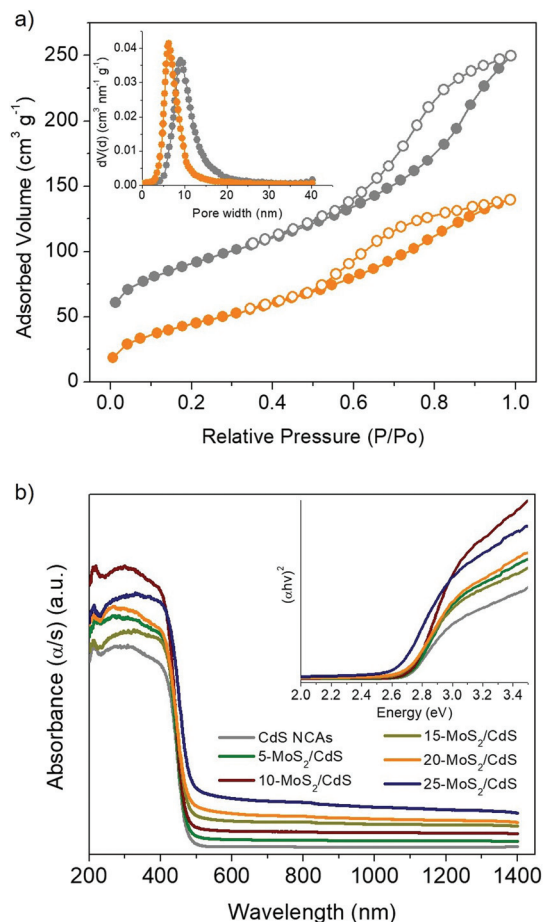


Fig. 3 (a) N₂ adsorption–desorption isotherms of the mesoporous CdS (gray symbols) and 20-MoS₂/CdS (orange symbols) NCAs. The isotherm data of CdS NCAs are shifted by 30 cm³ g⁻¹ for clarity. Inset: the NLDFT pore size distribution plots calculated from the adsorption data. (b) UV–vis/NIR diffuse reflectance spectra of mesoporous CdS and MoS₂/CdS NCAs with 5, 10, 15, 20 and 25 wt% MoS₂ content. Inset: the corresponding Tauc plots for direct bandgap semiconductors.

Union of Pure and Applied Chemistry classification), recommending mesoporous solids with interconnected pores.³⁸ The resulting mesoporous MoS₂/CdS architectures exhibit Brunauer–Emmett–Teller (BET) surface areas of 159–196 m² g⁻¹ and pore volumes of 0.20–0.26 cm³ g⁻¹, which are slightly smaller than the values of the CdS pristine sample (225 m² g⁻¹ and 0.28 cm³ g⁻¹, respectively). The pore width in these materials was estimated by applying the non-local density functional theory (NLDFT) fitting model on the adsorption data.³⁹ The results reveal that all the samples have a quite narrow distribution of pore sizes with a peak maximum at ~9.1 nm for CdS NCAs and ~6.1–8.1 nm for the MoS₂-modified catalysts (insets of Fig. 3a and S3†). The slight decrease in the porosity of composite materials with increasing MoS₂ content can be attributed to the successful deposition of MoS₂ NSs on the pore walls of mesoporous CdS, which is consistent with our TEM observations. The textural parameters for all the prepared materials are summarized in Table 1.

To determine the optical absorption properties of CdS and MoS₂/CdS NCAs, ultraviolet–visible/near-IR (UV–vis/NIR) diffuse reflectance spectroscopy was performed. As shown in Fig. 3b, the mesoporous CdS NCAs show an absorption edge of ~457 nm (2.71 eV) due to the intrinsic electron transition from the valence band (VB) to the CB in CdS NCs. Note that, compared with typical bulk CdS (it has a bandgap of 2.4 eV), CdS NCAs exhibit a notable blue shift in the interband absorption, which probably results from the quantum confinement effect arising from the very small size of the constituent CdS NCs (*ca.* 4–5 nm in size according to TEM observations). Meanwhile, the absorption edge position of the MoS₂-modified materials undergoes a small downshift upon increasing the concentration of MoS₂. In particular, the bandgap energy (E_g) of the MoS₂/CdS NCAs estimated according to the Tauc method [$(\alpha h\nu)^2 \propto (h\nu - E_g)$, where α is the absorption coefficient and $h\nu$ is the energy of incident photons] falls in the range of 2.63 to 2.71 eV, see the inset of Fig. 3b and Table 1. The tiny, yet noteworthy, small reduction of the energy bandgap is ascribed to the strong interactions between the CdS NCs and MoS₂ layers, which allows electron transport across the interface to take place. Moreover, the MoS₂-modified samples also showed a significant absorption tail in the 480–1400 nm range, which is most likely due to the visible and NIR light response of MoS₂ NSs. Indeed, the intensity of this absorption progressively increases with increasing MoS₂ content in composite samples and also correlates well with the absorption spectrum of MoS₂ (Fig. S4†). The MoS₂ NSs exhibit an energy gap of 1.62 eV that is greater than the typical bandgap of bulk MoS₂ (~1.3 eV), probably due to the small size and few-layered structure of the as-synthesized MoS₂ nanolayers.

3.2 Photocatalytic evolution of H₂

The photocatalytic H₂ evolution activities of the different nanocatalysts were investigated under visible light irradiation ($\lambda \geq 420$ nm), using an aqueous solution containing Na₂S/Na₂SO₃ as the sacrificial reagent. Fig. 4a compares the rates of H₂ production over the different MoS₂-modified catalysts; the H₂ evolution rates were determined during the first 2 h of illumination. It can be seen that bare CdS NCAs exhibit relatively poor H₂ production performance (0.06 mmol h⁻¹), which is primarily related to the fast recombination of photoexcited electron–hole pairs. On the contrary, MoS₂ modification of catalysts led to a significant increase in photocatalytic activity, which is by a factor of 6.3× compared to CdS NCAs. In particular, the H₂ evolution activity scales almost linearly with the MoS₂ content and reaches a maximum (0.39 mmol h⁻¹) at 20 wt%. The deposition of an excessive amount of MoS₂ (25 wt%) apparently results in the light shading effect on CdS nanoparticles and/or the formation of interfacial charge-carrier recombination centers, leading to a decrease in the H₂ production rate (0.29 mmol h⁻¹). It should be noted that when MoS₂ NSs were used as catalysts, no evolution of H₂ was observed under the examined conditions (results not shown). Also, blank experiments in the absence of light or catalyst did not show any evol-



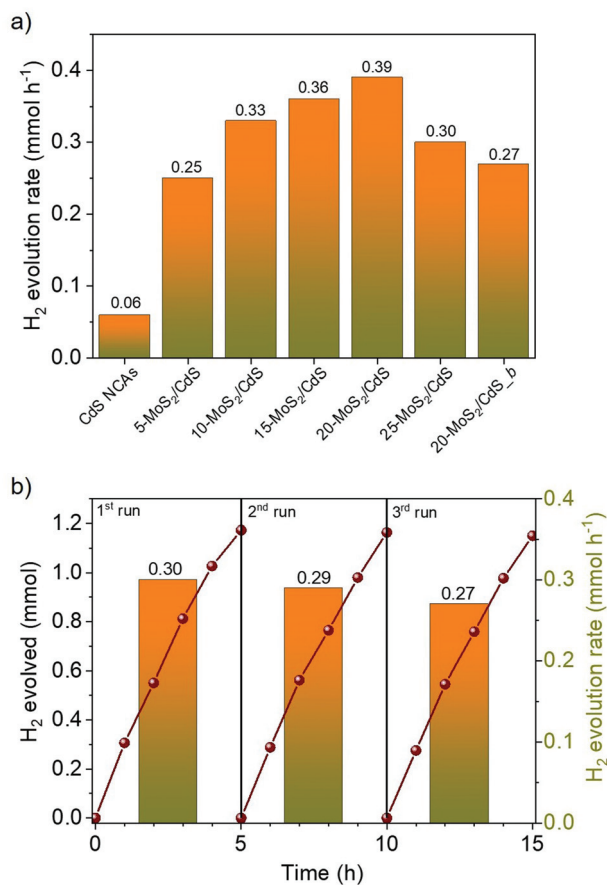


Fig. 4 (a) H₂ production rates for the mesoporous CdS and MoS₂/CdS NCAs. The H₂ evolution rate of the catalyst with 20 wt% bulk MoS₂ (20-MoS₂/CdS_b) is also given. (b) Photocatalytic recycling tests of the 20-MoS₂/CdS NCAs. The columns show the H₂ production rates of the 5 h time intervals. Reaction conditions: 20 mg catalyst (or 30 mg for recycling study), 20 mL aqueous solution of 0.35 M Na₂S and 0.25 M Na₂SO₃, 300 W Xenon light source with a cut-off filter ($\lambda \geq 420$ nm), 20 \pm 2 °C.

ution of H₂, indicating that H₂ is produced by a photocatalytic reaction.

Besides chemical composition, morphological effects are an important aspect of the efficiency of the MoS₂/CdS nanocatalysts. To elucidate this, we also prepared a reference catalyst by depositing 20 wt% of bulk MoS₂ flakes (*ca.* 1.2–3 μ m in lateral size)³¹ on the surface of mesoporous CdS NCAs (designated as 20-MoS₂/CdS_b), and their H₂ evolution performance was assessed under identical conditions (*i.e.*, using 20 mg of the 20-MoS₂/CdS_b catalyst dispersed in 0.35 M Na₂S and 0.25 M Na₂SO₃ aqueous solution; $\lambda \geq 420$ nm light radiation, 20 \pm 2 °C). EDS analysis indicated that this catalyst contains 21.5 wt% MoS₂ (Fig. S5a†). Also, it possesses almost identical textural and optical characteristics to those of the nanostructured 20-MoS₂/CdS analogue, as inferred by XRD, N₂ physisorption and UV-vis/NIR measurements, see Figs. S5b–d† and Table 1. Interestingly, the resultant 20-MoS₂/CdS_b catalyst showed a 1.4 \times decrease of the H₂ evolution performance

(0.27 mmol h⁻¹) over that of the 20-MoS₂/CdS nanocatalyst (Fig. 4a), despite exhibiting very similar chemical composition, porosity and light absorption abilities. Therefore, the pronounced enhancement in the photocatalytic performance of 20-MoS₂/CdS NCAs is clearly associated with the nanoscale interfacial contact between MoS₂ nanolayers and CdS NCs, which probably contributes to an increased concentration of exposed S-edge sites of MoS₂ and better electronic communication within the heterostructure. In general, reduction of the MoS₂ layer size could increase the number of bridging S₂²⁻ and terminal S²⁻ edge atoms, which are the catalytic active sites in MoS₂.⁴⁰ As for the facile electron transfer from CdS to MoS₂, this is further supported by photoluminescence and electrochemical studies (see below).

To optimize the reaction parameters, various photocatalytic H₂ evolution tests were carried out using different sacrificial hole scavengers and mass loadings of the 20-MoS₂/CdS catalyst. To this end, a series of sacrificial reagents, namely phenol, methanol, triethanolamine, triethylamine, lactic acid and Na₂S/Na₂SO₃, were examined in 20 mL of water that contained a fixed amount of catalyst. As shown in Fig. S6,† we obtained optimal results with the S²⁻/SO³⁻ mixture and used this sacrificial reagent during further studies. In general, S²⁻ and SO³⁻ ions not only enable H₂ production by consuming the surface-reaching holes, but also S²⁻ can replenish sulfur defects in metal sulfide materials originating from anodic photocorrosion.⁴¹ Also, photocatalytic tests using different concentrations of the 20-MoS₂/CdS catalyst indicated that the rate of H₂ evolution increases with increasing catalyst dose up to 1 g L⁻¹ (Fig. S7†). We interpret this as the saturation in light absorption by the catalyst particles. For higher catalyst concentrations (1.5 g L⁻¹), the slight reduction in the H₂ production rate (0.30 mmol h⁻¹) can be attributed to the light scattering effect of the catalyst particles. Overall, under the optimum reaction conditions (1 g L⁻¹ catalyst dose; Na₂S/Na₂SO₃ aqueous solution), 20-MoS₂/CdS NCAs catalyze water splitting with a H₂ production rate of \sim 0.4 mmol h⁻¹ (or \sim 19 mmol h⁻¹ g⁻¹ mass activity) and a calculated apparent quantum yield (AQY) of 51.2% at 420 \pm 10 nm. To the best of our knowledge, this efficiency far exceeds that of other MoS₂-modified CdS photocatalysts in previous reports. A comparison of the H₂ evolution activities of various MoS₂-modified CdS photocatalysts is presented in Table S1.†

Except for high photocatalytic activity, the mesoporous 20-MoS₂/CdS NCAs also exhibited persistent stability under the studied conditions. The stability of the catalyst was evaluated by repeated catalytic measurements for three cycles. Between each run, the catalyst was collected by centrifugation and re-suspended in a fresh Na₂S/Na₂SO₃ solution. Also, before each catalytic test, the reaction mixture was bubbled with Ar gas for at least 30 min, so that no H₂ and O₂ gases were detected by gas-chromatography (GC). As shown in Fig. 4b, 20-MoS₂/CdS presents a quite stable H₂ evolution rate, retaining approximately 90% of its initial activity after three runs. Nevertheless, a small decline of the H₂ production rate during the third run may be caused by minor photocorrosion and/or mass loss of



catalyst during the recycling tests. EDS and N₂ physisorption measurements were performed to verify the stability of the reused catalyst. The EDS spectra indicated a Mo/Cd atomic ratio of 19.9 : 80.1 that corresponds to a MoS₂ content of about 21.6 wt%, while N₂ physisorption data showed a surface area of 156 m² g⁻¹, pore size ~6 nm and pore volume of 0.16 cm³ g⁻¹ (Fig. S8†), very close to the values obtained from the fresh catalyst. Moreover, no change in the chemical state of elements in 20-MoS₂/CdS NCAs was observed after the stability test, as shown by XPS spectra (see Fig. S9†), substantiating the excellent stability of the catalyst during the photocatalytic reactions.

3.3 Role of MoS₂ NSs in the photochemical activity

To rationalize the role of MoS₂ NSs on the catalytic performance and to better understand the charge-transfer dynamics at the nanoscale MoS₂/CdS interfaces, we used electrochemical impedance spectroscopy (EIS). Fig. 5a depicts Mott–Schottky plots ($1/C_{sc}^2$ versus applied voltage E curves) of the mesoporous CdS and MoS₂/CdS NCAs; all the examined materials were drop-cast as thin films on a fluorinated tin oxide (FTO) glass substrate. From these plots, the flat band potentials (E_{FB}) of the samples were determined using the tangent lines of the $1/C_{sc}^2 - E$ curves. As shown in Table 2, the obtained E_{FB} values of MoS₂/CdS NCA catalysts range from -0.68 V to -0.75 V, whereas the E_{FB} position of CdS NCAs locates at -0.84 V; all

Table 2 Electrochemical results obtained from EIS analysis for MoS₂ NSs and different mesoporous MoS₂-modified CdS catalysts

Catalyst	E_{FB} (V vs. RHE, pH 7)	E_{VB}	Donor density (N_D , cm ⁻³)	R_{ct} (kΩ)
MoS ₂ NSs	-0.68	0.94	6.89×10^{17}	
CdS NCAs	-0.84	1.87	5.87×10^{16}	13.5
5-MoS ₂ /CdS	-0.75	1.96	1.17×10^{17}	6.3
10-MoS ₂ /CdS	-0.72	2.00	1.22×10^{17}	5.6
15-MoS ₂ /CdS	-0.71	1.97	1.43×10^{17}	5.3
20-MoS ₂ /CdS	-0.68	2.00	1.70×10^{17}	2.7
25-MoS ₂ /CdS	-0.69	1.94	1.65×10^{17}	4.5
20-MoS ₂ /CdS _b	-0.75	1.92	1.13×10^{17}	5.2

electrochemical potentials are reported versus the reversible hydrogen electrode (RHE) at pH 7. Moreover, the positive slopes of the $1/C_{sc}^2 - E$ plots express that electrons are the majority charge-carriers (n-type conductivity) in these materials. By combining the E_{FB} potentials with the respective bandgaps (as obtained from optical UV-vis/NIR spectroscopy), we obtained the band-edge potentials for each catalyst and the respective band-edge diagrams are presented in Fig. 5b. More specifically, for the calculation of band diagrams we have considered that the E_{FB} level is located very close to the CB minimum of heavily doped n-type semiconductors and thus

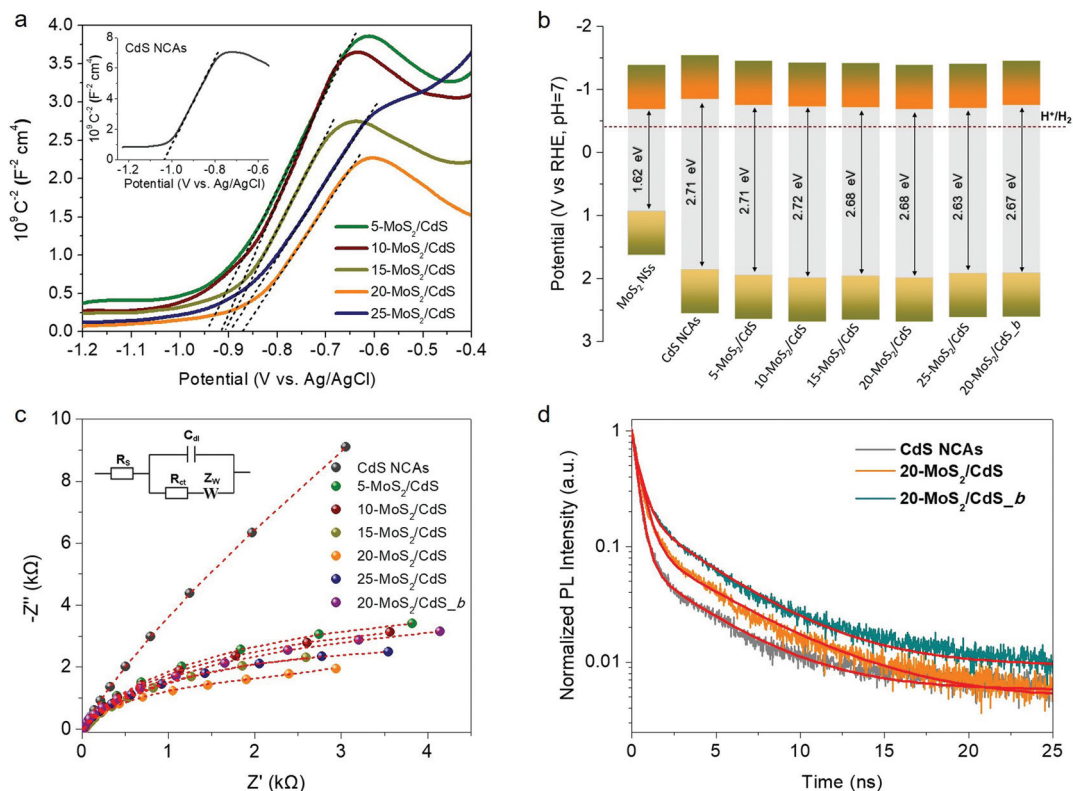


Fig. 5 (a) Mott–Schottky plots of the MoS₂-modified CdS NCAs (inset: the Mott–Schottky plot of CdS NCAs). (b) Band-edge positions and (c) Nyquist plots for MoS₂ NSs, CdS NCAs and various MoS₂-modified CdS NCAs (inset: equivalent circuit model $R_s[C_{dl}/(R_{ct} + Z_w)]$). (d) Comparative time-resolved photoluminescence (TRPL) decay profiles of pristine CdS and MoS₂-modified CdS NCAs with 20 wt% MoS₂ NSs (20-MoS₂/CdS) and MoS₂ bulk flakes (20-MoS₂/CdS_b).



the VB potential (E_{VB}) was obtained by adding the energy gap (E_g) to the E_{FB} level; the calculated E_{VB} values for each catalyst are listed in Table 2. As seen from the band diagrams, the E_{FB} level progressively undergoes an anodic shift from -0.84 V to -0.68 V with increasing the MoS₂ NS content on the surface of CdS. Moreover, as deduced from the magnitude of the slopes of $1/C_{SC}^2$ vs. E plots, MoS₂ modification also increases the donor density (N_D) of the catalysts. The measured N_D values for the MoS₂/CdS NCAs range from 1.17 to 1.70×10^{17} cm⁻³, which are significantly larger than that obtained for the unmodified CdS NCAs (5.87×10^{16} cm⁻³), as shown in Table 2. The observed anodic shift at the E_{FB} position of MoS₂/CdS NCAs can be interpreted by the lower E_{FB} of MoS₂ NSs (*i.e.*, -0.68 V vs. RHE, as determined through Mott–Schottky measurements, see Fig. S10a†) compared to that of CdS mesoporous. This results in the formation of an internal electric field across the MoS₂/CdS junction (upon contact) that drives electrons from CdS to MoS₂ until the Fermi levels reach equilibrium; for heavily n-doped semiconductors (such as CdS NCs and MoS₂ NSs), we assume that the Fermi level is located near to the CB edge position. Such an electron flow forms a depletion layer at the CdS surface, which lowers the CB energy potential closer to that of MoS₂. Indeed, the intrinsic electric field at the contact interface can result in an increase in charge separation and, therefore, in electron donor density (N_D) at the MoS₂/CdS junctions, which is in line with the above Mott–Schottky analysis results.

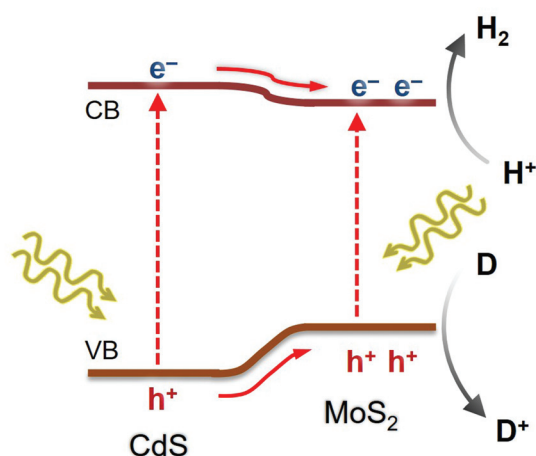
Moreover, the charge transfer properties of the prepared catalysts were also studied by EIS Nyquist plots. Fig. 5c shows the Nyquist plots of pristine and MoS₂-modified CdS samples recorded in the 0.5 M Na₂SO₄ electrolyte under open circuit conditions over a frequency span from 1 Hz to 1 MHz. A typical Randles equivalent circuit was used to fit the experimental data consisting of a solution resistance (R_s), a charge-transfer resistance (R_{ct}), a double layer capacitance (C_{dl}) and a diffusional resistance element (Z_w) (see the inset of Fig. 5c), and the results of the fitting parameters are given in Table S2.† Fitting results indicated that the MoS₂-modified catalysts have R_{ct} values of ~ 2.7 – 6.3 k Ω , which are significantly lower than that of CdS NCAs (~ 13.5 k Ω) (Table 2), suggesting a more favorable charge transfer across the MoS₂/CdS junctions. Subsequently, this process contributes to the improved photocatalytic efficiency of the MoS₂-modified catalysts. It is worth mentioning that among diverse MoS₂/CdS catalysts, 20-MoS₂/CdS NCAs possessed the lowest resistance in carrier mobility (~ 2.7 k Ω), which associates well with their remarkable photocatalytic H₂ evolution performance. Notably, the bulk reference catalyst (20-MoS₂/CdS_b) manifests a relatively higher R_{ct} value (~ 5.2 k Ω) than the nanostructured analogue (20-MoS₂/CdS), thereby suggesting more sluggish charge transfer kinetics, despite having similar composition and band edge positions (see Fig. 5b and Fig. S10b†). In addition, as shown in Table 2, the charge carrier density of 20-MoS₂/CdS_b is lower than that of 20-MoS₂/CdS according to the Mott–Schottky equation (1.13×10^{17} vs. 1.70×10^{17} cm⁻³), suggesting an inferior electronic contact between the bulk MoS₂ layers and CdS nanostructure.

This observation agrees with Nyquist charge-transfer resistance measurements and confirms that the nanoscale dimensionality of the MoS₂ layers has a prominent impact on the charge separation dynamics at the MoS₂/CdS interfaces. This is also elaborated by the photoluminescence (PL) study of catalysts under 375 nm excitation wavelength. As shown in Fig. S11,† the 20-MoS₂/CdS catalyst shows a weaker PL emission at ~ 460 nm (corresponding to the near band-edge excitonic relaxation) than the pristine CdS and bulk 20-MoS₂/CdS_b samples, indicating better mitigation of electron–hole recombination by the nanoscale MoS₂/CdS junctions.

The dynamics of charge-carrier recombination in these materials was also investigated using time-resolved photoluminescence (TRPL) decay measurements upon monochromatic excitation at 375 nm. To extract the PL lifetimes, we used a biexponential function [$F(t) = y_0 + \sum_i \alpha_i e^{-t/\tau_i}$, $i = 1, 2$] to fit the PL decay data, where α_i denotes the amplitude fractions ($\sum_i \alpha_i = 1$) and τ_i is the carrier lifetime (Fig. 5d); we used a two-component equation describing the surface and possible defect-mediated (fast) and bulk (slow) radiative charge-carrier recombination, respectively. Through this fitting analysis the average lifetimes (τ_{av}) of mesoporous CdS, 20-MoS₂/CdS and 20-MoS₂/CdS_b catalysts were determined as 3.12, 4.15 and 3.46 ns, respectively; the fitting parameters of the PL decay curves are listed in Table S3.† Once again, TRPL results evidence that the formation of MoS₂/CdS nanojunctions has a prominent impact on the charge-transfer kinetics at the catalyst interface, promoting a better separation of charge carriers and a more efficient utilization of the long-lived photoexcited electrons for the H₂ evolution reaction, in line with the respective decrease of PL intensity shown in Fig. S11.† This is also apparent in the current density–voltage (J – V) data provided in Fig. S12,† which indicates a substantially lower proton reduction potential (by 0.18 V) in MoS₂-modified catalysts compared to the pristine CdS. The observed positive shift in the onset potential with MoS₂ deposition can be ascribed to the increased charge separation at the MoS₂/CdS interface and the lower energy barrier for H₂ evolution at the MoS₂ edge sites. Moreover, in comparison to the bulk reference catalyst (20-MoS₂/CdS_b), 20-MoS₂/CdS NCAs exhibited a quick current response, implying more rapid reaction kinetics caused by the low dimensionality of MoS₂ layers and increased amount of active MoS₂ edge-sites. For example, the current density at an applied voltage of -1.1 V versus Ag/AgCl is greatly enhanced from 0.65 mA cm⁻² to 1.34 mA cm⁻² with 20-MoS₂/CdS_b and 20-MoS₂/CdS catalyst, respectively.

According to the above analysis, a possible reaction mechanism for the photocatalytic H₂ evolution reaction by MoS₂/CdS nanocatalysts was suggested, as presented in Scheme 1. Light irradiation first excites the electrons from the VB of CdS and MoS₂ to their respective CB, producing photogenerated carriers. Subsequently, due to the lower CB potential of MoS₂ than CdS and the intrinsic electric field formed at their contact interface, electrons are transferred from CdS to the





Scheme 1 Expected charge transfer during photocatalytic H₂ production on mesoporous MoS₂/CdS NCAs under visible light excitation.

active edge sites of MoS₂ NSs, where they efficiently reduce H⁺ to H₂. The smooth charge transfer and abated electron–hole recombination, which are effectively induced by the nanoscale MoS₂/CdS junctions, would provide prolonged lifetime and thus more possibilities for electrons to engage in catalytic proton reduction. In parallel, the photoexcited holes on the surface of CdS and/or MoS₂ oxidize the sacrificial reagents (such as S²⁻ and SO₃²⁻). Evidence of such charge-transfer pathways within the MoS₂/CdS nanojunction network was found from EIS and TRPL studies.

4. Conclusions

In summary, nanojunction networks of sub 5 nm sized CdS NCs and ultrasmall MoS₂ NSs (*ca.* 10–15 nm in lateral size) with different compositions have been successfully synthesized by depositing MoS₂ NSs on the surface of mesoporous CdS NC assemblies. The chemical composition, crystallinity and morphology of the prepared catalysts were confirmed by electron microprobe analysis, X-ray diffraction, electron microscopy, N₂ porosimetry and X-ray photoelectron spectroscopy measurements. These mesoporous structures have a proper electronic structure that allows spatial separation and transport of charge carriers across the MoS₂/CdS nano-interfaces and promotes high H₂ production efficiency of the photoexcited electrons. Evidence for this was provided by photoluminescence and electrochemical impedance spectroscopy. Owing to these unique characteristics, the MoS₂/CdS nanojunction network with the optimized MoS₂ content (20 wt%) attains a 6.3× and 1.4× increase in the H₂ production rate compared to bare mesoporous CdS and the bulk MoS₂/CdS analogue, respectively. It also presents an energy conversion efficiency of 51.2% at 420 ± 10 nm, which is one of the highest reported for MoS₂-modified CdS catalysts. The recycling test indicated persistent H₂ evolution activity and excellent chemical and structural stability of the nanoparticle-assembled structure. The results

of this study provide an understanding of the charge transfer dynamics at the MoS₂/CdS nanoscale junctions and support the potential viability of the present MoS₂-modified CdS nano-architectures as photocatalysts for clean energy conversion.

Author contributions

Ch. Patriarchea: investigation, formal analysis, writing – original draft; I. Vamvasakis: investigation, formal analysis, writing – original draft; E. D. Koutsouroubi: investigation, formal analysis; G. S. Armatas: conceptualization, supervision, resources, writing – review and editing.

Conflicts of interest

There are no conflicts to declare.

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