

## RESEARCH ARTICLE

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# Enhancing interfacial charge transfer in mesoporous MoS<sub>2</sub>/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 MoS<sub>2</sub> 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 MoS<sub>2</sub> 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 N<sub>2</sub> physisorption prove that the MoS<sub>2</sub>-modified CdS nanocatalysts are composed of a porous network of connected cubic CdS nanocrystals and 2H-phase MoS<sub>2</sub> nanosheets and possess a high internal BET surface area (ca. 159–225 m<sup>2</sup> g<sup>-1</sup>) 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 MoS<sub>2</sub>/CdS junctions provide more efficient electronic connectivity and charge carrier dissociation across the catalyst interfaces, resulting in a remarkable enhancement in the photocatalytic H<sub>2</sub> production activity. The optimized MoS<sub>2</sub>/CdS catalyst at 20 wt% MoS<sub>2</sub> content achieves a H<sub>2</sub> production rate up to ~0.4 mmol h<sup>-1</sup> (or ~19 mmol h<sup>-1</sup> g<sup>-1</sup> mass activity) with remarkable stability under visible light irradiation, corresponding to an overall 6.7x enhancement of H<sub>2</sub> 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.<sup>1,2</sup> 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.<sup>3</sup> Thus far, a great variety of semiconducting materials, including metal oxides<sup>4</sup> and sulfides,<sup>5,6</sup> 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.<sup>7–9</sup> 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<sup>+</sup> reduction potential (–0.41 V, pH 7).<sup>10–12</sup> 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.<sup>13,14</sup> 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.<sup>15,16</sup>

After the revolution of graphene, transition metal dichalcogenides (TMDs), with the formula  $\text{MX}_2$  where M is Mo, W, etc. and X is S, Se, and Te, have also received attention as ideal co-catalysts for the photocatalytic  $\text{H}_2$  evolution reaction.<sup>17</sup> Among TMDs, 2D molybdenum disulfide ( $\text{MoS}_2$ ), 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.<sup>18</sup> In the bulk or multilayer form,  $\text{MoS}_2$  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),  $\text{MoS}_2$  also adopts two metastable phases, that is, the rhombohedral 3R (semiconducting phase) and trigonal 1T (metallic phase) structure.<sup>19</sup> Among these polytypes, 2H- $\text{MoS}_2$  although exhibits lower mobility for transporting charge carriers than the metallic 1T- $\text{MoS}_2$ , shows better stability and interesting semiconducting properties that are useful for applications in light energy conversion and optoelectronics.<sup>20</sup> Also, the 2H phase of  $\text{MoS}_2$  can exhibit better photocatalytic activities than the semiconducting 3R counterpart due to the unexfoliated structure of 3R- $\text{MoS}_2$ . In particular, the three-layered (3R)  $\text{MoS}_2$  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.<sup>21</sup> For this purpose, considering the semiconducting properties of  $\text{MoS}_2$ , there have been extensive studies on various heterostructures using 2H- $\text{MoS}_2$  as the co-catalyst for improved  $\text{H}_2$  production yield.<sup>22,23</sup> For instance, Zhang *et al.* demonstrated the selective growth of single-layer  $\text{MoS}_2$  nanoflakes, with a 4–10 nm lateral size, on the Cd-rich (0001) surface of wurtzite CdS particles.<sup>24</sup> This catalyst was reported to undergo photocatalytic reduction of water in the presence of lactic acid as the hole scavenger, yielding a  $\text{H}_2$  evolution rate of  $1.47 \text{ mmol h}^{-1} \text{ g}^{-1}$ . Song's group successfully prepared a 2D/2D heterojunction by assembling ultrathin  $\text{MoS}_2$  sheets and exfoliated CdS nanolayers through an adsorption–calcination process.<sup>25</sup> They found that the optimal photocatalyst presented a maximum photocatalytic  $\text{H}_2$  production activity of  $18.43 \text{ mmol h}^{-1} \text{ g}^{-1}$ . More recently, Fujishima and coworkers coupled 2D  $\text{MoS}_2$  thin sheets as  $\text{H}_2$  evolution catalysts with CdS microspheres by a template-free solvothermal method.<sup>26</sup> The nanostructured core–shell CdS@ $\text{MoS}_2$  particles showed high carrier transport efficiency and a remarkable  $\text{H}_2$  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.<sup>27</sup> 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.<sup>28</sup> 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- $\text{MoS}_2$  nanosheets (NSs) with an ultrasmall lateral size (*ca.* 10–15 nm) and evaluate their performance as catalysts for water splitting and  $\text{H}_2$  evolution under visible light. The 2D/3D  $\text{MoS}_2$ /CdS nano-heterostructures with various loadings of  $\text{MoS}_2$  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  $\text{MoS}_2$ /CdS contacts minimizes electron–hole recombination and increases the electron density and mobility at the  $\text{MoS}_2$ /CdS junction, yielding accelerated water reduction kinetics. Owing to these characteristics, the optimized catalyst with 20 wt%  $\text{MoS}_2$  NSs exhibits a significant enhancement of the photocatalytic activity, yielding an apparent quantum yield (AQY) of 51.2% for  $\text{H}_2$  evolution at 420 nm, a record efficiency for  $\text{MoS}_2$ -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,  $\text{CdCl}_2$  (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  $\text{NH}_4\text{OH}$ . Next, a  $\text{Na}_2\text{S}$  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.<sup>27</sup> 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  $\text{NH}_4\text{OH}$  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,  $\text{H}_2\text{O}_2$  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 MoS<sub>2</sub> NS synthesis.** For the synthesis of the 2H-MoS<sub>2</sub> NSs, monovacant Keggin-type phosphomolybdate (Na<sub>7</sub>[PMo<sub>11</sub>O<sub>39</sub>]) was synthesized following a previously reported procedure.<sup>30</sup> Next, Na<sub>7</sub>[PMo<sub>11</sub>O<sub>39</sub>] (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, MoS<sub>2</sub> microflakes were also synthesized following a similar procedure, but using (NH<sub>4</sub>)<sub>6</sub>Mo<sub>7</sub>O<sub>24</sub>·4H<sub>2</sub>O (0.0275 mmol) as the Mo source.

**2.1.4 Mesoporous MoS<sub>2</sub>/CdS NCAs.** Porous structures of MoS<sub>2</sub> NSs and CdS NCs with different MoS<sub>2</sub> contents, namely, 5, 10, 15, 20 and 25 wt%, were prepared by a facile deposition method. For a typical synthesis of a 20 wt% MoS<sub>2</sub>-modified CdS sample, the as-prepared MoS<sub>2</sub> NSs (20 mg) were added to a H<sub>2</sub>O : 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 MoS<sub>2</sub> NSs was formed. In a separate vial, mesoporous CdS NCAs (80 mg) were dispersed in a H<sub>2</sub>O : IPA solution (2 : 1 v/v, 5 mL) under vigorous stirring at RT. Finally, the suspension of exfoliated MoS<sub>2</sub> 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% MoS<sub>2</sub>-modified CdS samples were synthesized following a similar procedure using the stoichiometric amount of MoS<sub>2</sub> NSs for each case.

A bulk-like MoS<sub>2</sub>-modified CdS catalyst (designated as 20-MoS<sub>2</sub>/CdS<sub>b</sub>) was also prepared by depositing 20 wt% of MoS<sub>2</sub> microflakes on the surface of mesoporous CdS.

## 2.2 Photocatalytic water splitting study

The photocatalytic H<sub>2</sub> 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 Na<sub>2</sub>SO<sub>3</sub> and 0.35 M Na<sub>2</sub>S (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  $\mu$ L of gas was collected from the reactor's headspace and the generated H<sub>2</sub> 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 $\alpha$  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 $\alpha$  and S-K $\alpha$  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 (LaB<sub>6</sub> 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 $\alpha$  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 BaSO<sub>4</sub> 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  $\alpha$  and  $S$  are the absorption and scattering coefficients, respectively] into absorption spectra. N<sub>2</sub> 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<sup>–5</sup> 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 N<sub>2</sub> 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 Na<sub>2</sub>SO<sub>4</sub> 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  $\mu$ 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  $\mu$ L of the solution was dropped on a fluorinated tin oxide (FTO, 10  $\Omega$  sq<sup>–1</sup>) substrate (effective



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

The space-charge capacitance ( $C_{\text{sc}}$ ) 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  $E_{\text{RHE}}$  is the potential in the RHE scale and  $E_{\text{Ag/AgCl}}$  is the measured potential in the Ag/AgCl scale.

The donor density ( $N_{\text{D}}$ ) 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,  $C_{\text{sc}}$  is the space charge capacitance,  $E_{\text{FB}}$  is the flat band potential,  $E$  is the applied potential,  $N_{\text{D}}$  is the donor density of the electrode material,  $\epsilon$  is the dielectric constant (8.9 for CdS),  $\epsilon_0$  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-MoS<sub>2</sub> NSs were produced using a polymer-templated oxidative aggregating method followed by a wet-chemical deposition process. Initially, 2D nanolayers of 2H-MoS<sub>2</sub> were prepared through a hydrothermal sulfurization reaction, using Na<sub>7</sub>[PMo<sub>11</sub>O<sub>39</sub>] and thioacetamide (CH<sub>3</sub>CSNH<sub>2</sub>) as the Mo and S source, respectively.<sup>31</sup> After annealing to a certain temperature, thioacetamide begins to decompose and evolve H<sub>2</sub>S that is capable of transforming MoO<sub>x</sub><sup>n-</sup> species into MoS<sub>2</sub>. The X-ray diffraction (XRD) pattern of as-made MoS<sub>2</sub> NSs, in Fig. S1,† indicates the successful chemical transformation of phosphomolybdate clusters into MoS<sub>2</sub> under our liquid-phase conditions; all the diffraction peaks can be assigned to the hexagonal (2H) structure of MoS<sub>2</sub>. Besides, the broadened XRD diffractions point to the nanoscale dimension of the MoS<sub>2</sub> 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. MoS<sub>2</sub> modification of the mesoporous CdS structure was realized by depositing the liquid exfoliated MoS<sub>2</sub> NSs onto the surface of CdS NCAs. Following this process, several *n*-MoS<sub>2</sub>/CdS NCAs with different compositions of grafted MoS<sub>2</sub>, 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 MoS<sub>2</sub>/CdS NCAs is very close to the nominal composition of the samples (within an 8% deviation), suggesting the complete binding of MoS<sub>2</sub> 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 MoS<sub>2</sub>/CdS NCAs display three Bragg diffraction peaks at  $2\theta$  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 MoS<sub>2</sub> deposition, CdS still maintains its crystallinity, indicating that MoS<sub>2</sub> nanolayers are mainly studded on the CdS surface. In XRD patterns, however, no additional peak is identified after MoS<sub>2</sub> deposition, even in the XRD plot of the composite sample with 25 wt% MoS<sub>2</sub>, which is due to the good dispersion and nanoscale size of the MoS<sub>2</sub> layers.

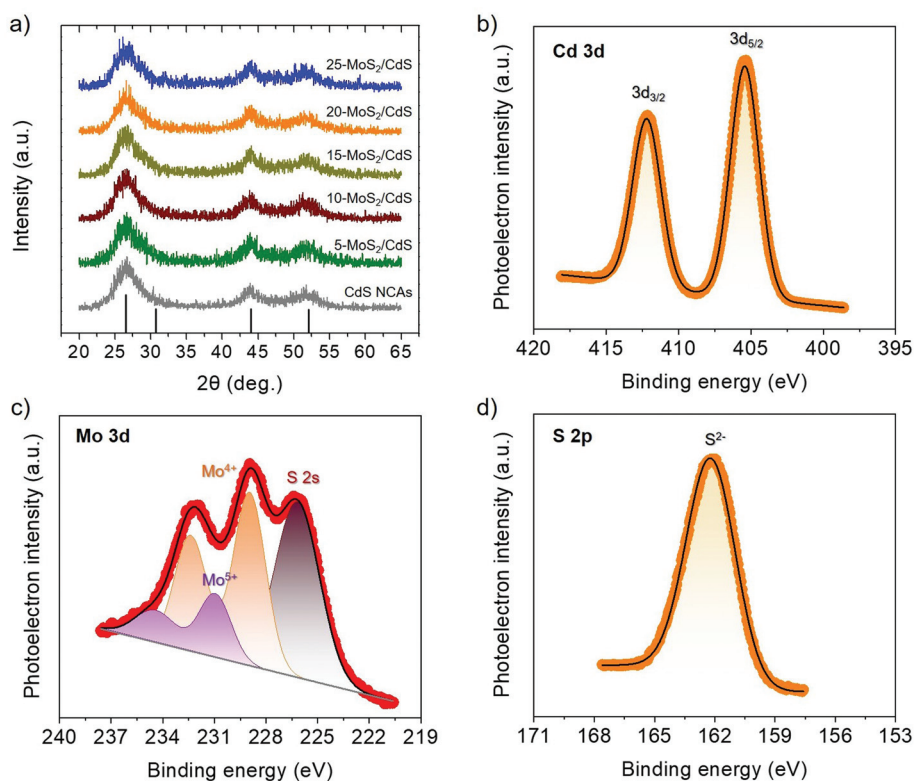
Typical XPS spectra of the best catalyst (20-MoS<sub>2</sub>/CdS) are shown in Fig. 1b–d. The XPS Cd 3d spectrum, in Fig. 1b, presents a doublet peak at  $405.3$  and  $412.2 \pm 0.2 \text{ eV}$  binding energies, corresponding to the Cd 3d<sub>5/2</sub> and Cd 3d<sub>3/2</sub> core level lines of Cd<sup>2+</sup> in CdS, respectively.<sup>32</sup> The XPS spectrum of the Mo 3d scan confirmed the presence of Mo<sup>4+</sup> species in MoS<sub>2</sub>, showing two peaks at  $229.1 \pm 0.2 \text{ eV}$  (3d<sub>5/2</sub>) and  $232.4 \pm 0.2 \text{ eV}$  (3d<sub>3/2</sub>) binding energies with a spin-orbit splitting of  $3.3 \text{ eV}$  (Fig. 1c), in line with other reports.<sup>33</sup> Besides the prominent Mo<sup>4+</sup> 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 MoS<sub>2</sub>/CdS NCAs

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

<sup>a</sup> Based on EDS analysis.





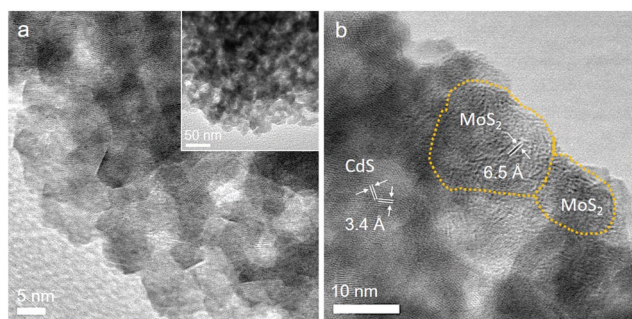
**Fig. 1** (a) XRD plots for the mesoporous CdS and MoS<sub>2</sub>/CdS NCAs. The standard diffraction data of the zinc blende CdS according to the JCPDS card no. 42-1411 are also shown (black bars). (b–d) Typical XPS core-level spectra of the Cd 3d (b), Mo 3d (c) and S 2p (d) region for the 20-MoS<sub>2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

can be assigned to unsaturated Mo<sup>4+</sup> atoms and/or higher oxidation state of Mo<sup>5+</sup> species due to mild surface oxidation after the synthesis process.<sup>34</sup> As for the singlet peak observed at  $226.6 \pm 0.2$  eV, this is associated with the S 2s signal of sulfides (S<sup>2-</sup>).<sup>35</sup> Furthermore, the XPS S 2p spectrum (Fig. 1d) exhibited a prominent peak at  $162.2 \pm 0.3$  eV that is characteristic of the S<sup>2-</sup> ions in metal sulfides.<sup>36</sup> Quantitative analysis of the XPS spectra also indicated a Mo-to-Cd atomic ratio close to 20.4 : 79.6 that corresponds to a  $\sim 22.1$  wt% MoS<sub>2</sub> 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-

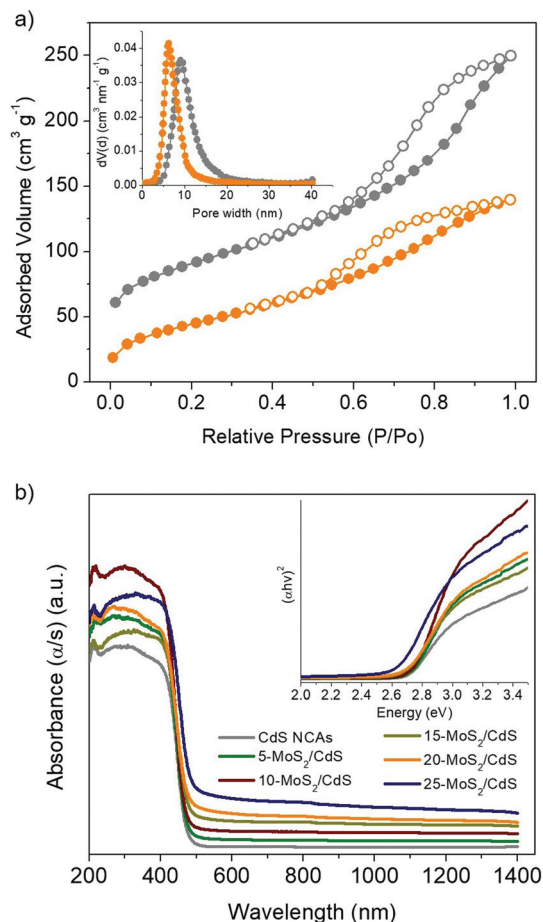
tive TEM images for the 20-MoS<sub>2</sub>/CdS sample are shown in Fig. 2. The images reveal that the framework of 20-MoS<sub>2</sub>/CdS is composed of approximately 4–5 nm sized CdS interconnected nanoparticles (Fig. 2a). In addition, from the high-resolution TEM (HRTEM) observation in Fig. 2b, some 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  $\sim 6.5$  Å that reasonably correlates to the (002) planes of MoS<sub>2</sub> NSs. This suggests a small enlargement of the interlayer distance of MoS<sub>2</sub> NSs with respect to the bulk MoS<sub>2</sub> (6.2 Å), which is due primarily to the small lateral size of the MoS<sub>2</sub> layers. A similar effect was previously observed for other MoS<sub>2</sub> nanostructures.<sup>37</sup> Taken together with XPS characterization of the surface structure, these results confirm that small-size MoS<sub>2</sub> 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 N<sub>2</sub> 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-MoS<sub>2</sub>/CdS NCAs are compared as shown in Fig. 3a to those of unmodified CdS NCAs. The respective plots for the other MoS<sub>2</sub>-modified catalysts (*i.e.*, with 5, 10, 15 and 25 wt% MoS<sub>2</sub> content) are depicted in Fig. S3.† All samples display characteristic type-IV isotherms with a H<sub>2</sub>-type hysteresis loop (according to the International



**Fig. 2** Representative (a) TEM and (b) HRTEM images of the 20-MoS<sub>2</sub>/CdS NCAs catalyst.





**Fig. 3** (a) N<sub>2</sub> adsorption–desorption isotherms of the mesoporous CdS (gray symbols) and 20-MoS<sub>2</sub>/CdS (orange symbols) NCAs. The isotherm data of CdS NCAs are shifted by 30 cm<sup>3</sup> g<sup>-1</sup> 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 MoS<sub>2</sub>/CdS NCAs with 5, 10, 15, 20 and 25 wt% MoS<sub>2</sub> content. Inset: the corresponding Tauc plots for direct bandgap semiconductors.

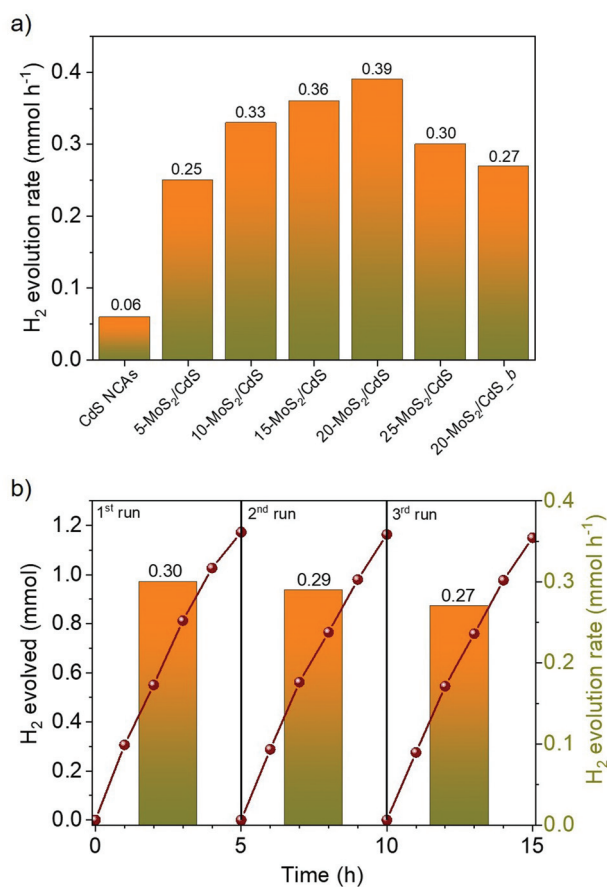
Union of Pure and Applied Chemistry classification), recommending mesoporous solids with interconnected pores.<sup>38</sup> The resulting mesoporous MoS<sub>2</sub>/CdS architectures exhibit Brunauer–Emmett–Teller (BET) surface areas of 159–196 m<sup>2</sup> g<sup>-1</sup> and pore volumes of 0.20–0.26 cm<sup>3</sup> g<sup>-1</sup>, which are slightly smaller than the values of the CdS pristine sample (225 m<sup>2</sup> g<sup>-1</sup> and 0.28 cm<sup>3</sup> g<sup>-1</sup>, respectively). The pore width in these materials was estimated by applying the non-local density functional theory (NLDFT) fitting model on the adsorption data.<sup>39</sup> 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 MoS<sub>2</sub>-modified catalysts (insets of Fig. 3a and S3†). The slight decrease in the porosity of composite materials with increasing MoS<sub>2</sub> content can be attributed to the successful deposition of MoS<sub>2</sub> 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 MoS<sub>2</sub>/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 MoS<sub>2</sub>-modified materials undergoes a small downshift upon increasing the concentration of MoS<sub>2</sub>. In particular, the bandgap energy ( $E_g$ ) of the MoS<sub>2</sub>/CdS NCAs estimated according to the Tauc method [ $(\alpha h\nu)^2 \propto (h\nu - E_g)$ , where,  $\alpha$  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 MoS<sub>2</sub> layers, which allows electron transport across the interface to take place. Moreover, the MoS<sub>2</sub>-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 MoS<sub>2</sub> NSs. Indeed, the intensity of this absorption progressively increases with increasing MoS<sub>2</sub> content in composite samples and also correlates well with the absorption spectrum of MoS<sub>2</sub> (Fig. S4†). The MoS<sub>2</sub> NSs exhibit an energy gap of 1.62 eV that is greater than the typical bandgap of bulk MoS<sub>2</sub> (~1.3 eV), probably due to the small size and few-layered structure of the as-synthesized MoS<sub>2</sub> nanolayers.

### 3.2 Photocatalytic evolution of H<sub>2</sub>

The photocatalytic H<sub>2</sub> evolution activities of the different nanocatalysts were investigated under visible light irradiation ( $\lambda \geq 420$  nm), using an aqueous solution containing Na<sub>2</sub>S/Na<sub>2</sub>SO<sub>3</sub> as the sacrificial reagent. Fig. 4a compares the rates of H<sub>2</sub> production over the different MoS<sub>2</sub>-modified catalysts; the H<sub>2</sub> evolution rates were determined during the first 2 h of illumination. It can be seen that bare CdS NCAs exhibit relatively poor H<sub>2</sub> production performance (0.06 mmol h<sup>-1</sup>), which is primarily related to the fast recombination of photoexcited electron–hole pairs. On the contrary, MoS<sub>2</sub> 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 H<sub>2</sub> evolution activity scales almost linearly with the MoS<sub>2</sub> content and reaches a maximum (0.39 mmol h<sup>-1</sup>) at 20 wt%. The deposition of an excessive amount of MoS<sub>2</sub> (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 H<sub>2</sub> production rate (0.29 mmol h<sup>-1</sup>). It should be noted that when MoS<sub>2</sub> NSs were used as catalysts, no evolution of H<sub>2</sub> 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) H<sub>2</sub> production rates for the mesoporous CdS and MoS<sub>2</sub>/CdS NCAs. The H<sub>2</sub> evolution rate of the catalyst with 20 wt% bulk MoS<sub>2</sub> (20-MoS<sub>2</sub>/CdS\_b) is also given. (b) Photocatalytic recycling tests of the 20-MoS<sub>2</sub>/CdS NCAs. The columns show the H<sub>2</sub> 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 Na<sub>2</sub>S and 0.25 M Na<sub>2</sub>SO<sub>3</sub>, 300 W Xenon light source with a cut-off filter ( $\lambda \geq 420$  nm), 20  $\pm$  2  $^{\circ}$ C.

ution of H<sub>2</sub>, indicating that H<sub>2</sub> is produced by a photocatalytic reaction.

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

(0.27 mmol h<sup>-1</sup>) over that of the 20-MoS<sub>2</sub>/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-MoS<sub>2</sub>/CdS NCAs is clearly associated with the nanoscale interfacial contact between MoS<sub>2</sub> nanolayers and CdS NCs, which probably contributes to an increased concentration of exposed S-edge sites of MoS<sub>2</sub> and better electronic communication within the heterostructure. In general, reduction of the MoS<sub>2</sub> layer size could increase the number of bridging S<sub>2</sub><sup>2-</sup> and terminal S<sup>2-</sup> edge atoms, which are the catalytic active sites in MoS<sub>2</sub>.<sup>40</sup> As for the facile electron transfer from CdS to MoS<sub>2</sub>, this is further supported by photoluminescence and electrochemical studies (see below).

To optimize the reaction parameters, various photocatalytic H<sub>2</sub> evolution tests were carried out using different sacrificial hole scavengers and mass loadings of the 20-MoS<sub>2</sub>/CdS catalyst. To this end, a series of sacrificial reagents, namely phenol, methanol, triethanolamine, triethylamine, lactic acid and Na<sub>2</sub>S/Na<sub>2</sub>SO<sub>3</sub>, were examined in 20 mL of water that contained a fixed amount of catalyst. As shown in Fig. S6,<sup>†</sup> we obtained optimal results with the S<sup>2-</sup>/SO<sub>3</sub><sup>3-</sup> mixture and used this sacrificial reagent during further studies. In general, S<sup>2-</sup> and SO<sub>3</sub><sup>3-</sup> ions not only enable H<sub>2</sub> production by consuming the surface-reaching holes, but also S<sup>2-</sup> can replenish sulfur defects in metal sulfide materials originating from anodic photocorrosion.<sup>41</sup> Also, photocatalytic tests using different concentrations of the 20-MoS<sub>2</sub>/CdS catalyst indicated that the rate of H<sub>2</sub> evolution increases with increasing catalyst dose up to 1 g L<sup>-1</sup> (Fig. S7<sup>†</sup>). We interpret this as the saturation in light absorption by the catalyst particles. For higher catalyst concentrations (1.5 g L<sup>-1</sup>), the slight reduction in the H<sub>2</sub> production rate (0.30 mmol h<sup>-1</sup>) can be attributed to the light scattering effect of the catalyst particles. Overall, under the optimum reaction conditions (1 g L<sup>-1</sup> catalyst dose; Na<sub>2</sub>S/Na<sub>2</sub>SO<sub>3</sub> aqueous solution), 20-MoS<sub>2</sub>/CdS NCAs catalyze water splitting with a H<sub>2</sub> production rate of  $\sim$ 0.4 mmol h<sup>-1</sup> (or  $\sim$ 19 mmol h<sup>-1</sup> g<sup>-1</sup> 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 MoS<sub>2</sub>-modified CdS photocatalysts in previous reports. A comparison of the H<sub>2</sub> evolution activities of various MoS<sub>2</sub>-modified CdS photocatalysts is presented in Table S1.<sup>†</sup>

Except for high photocatalytic activity, the mesoporous 20-MoS<sub>2</sub>/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 Na<sub>2</sub>S/Na<sub>2</sub>SO<sub>3</sub> solution. Also, before each catalytic test, the reaction mixture was bubbled with Ar gas for at least 30 min, so that no H<sub>2</sub> and O<sub>2</sub> gases were detected by gas-chromatography (GC). As shown in Fig. 4b, 20-MoS<sub>2</sub>/CdS presents a quite stable H<sub>2</sub> evolution rate, retaining approximately 90% of its initial activity after three runs. Nevertheless, a small decline of the H<sub>2</sub> production rate during the third run may be caused by minor photocorrosion and/or mass loss of



catalyst during the recycling tests. EDS and  $N_2$  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  $MoS_2$  content of about 21.6 wt%, while  $N_2$  physisorption data showed a surface area of  $156 \text{ m}^2 \text{ g}^{-1}$ , pore size  $\sim 6 \text{ nm}$  and pore volume of  $0.16 \text{ cm}^3 \text{ g}^{-1}$  (Fig. S8<sup>†</sup>), very close to the values obtained from the fresh catalyst. Moreover, no change in the chemical state of elements in 20- $MoS_2$ /CdS NCAs was observed after the stability test, as shown by XPS spectra (see Fig. S9<sup>†</sup>), substantiating the excellent stability of the catalyst during the photocatalytic reactions.

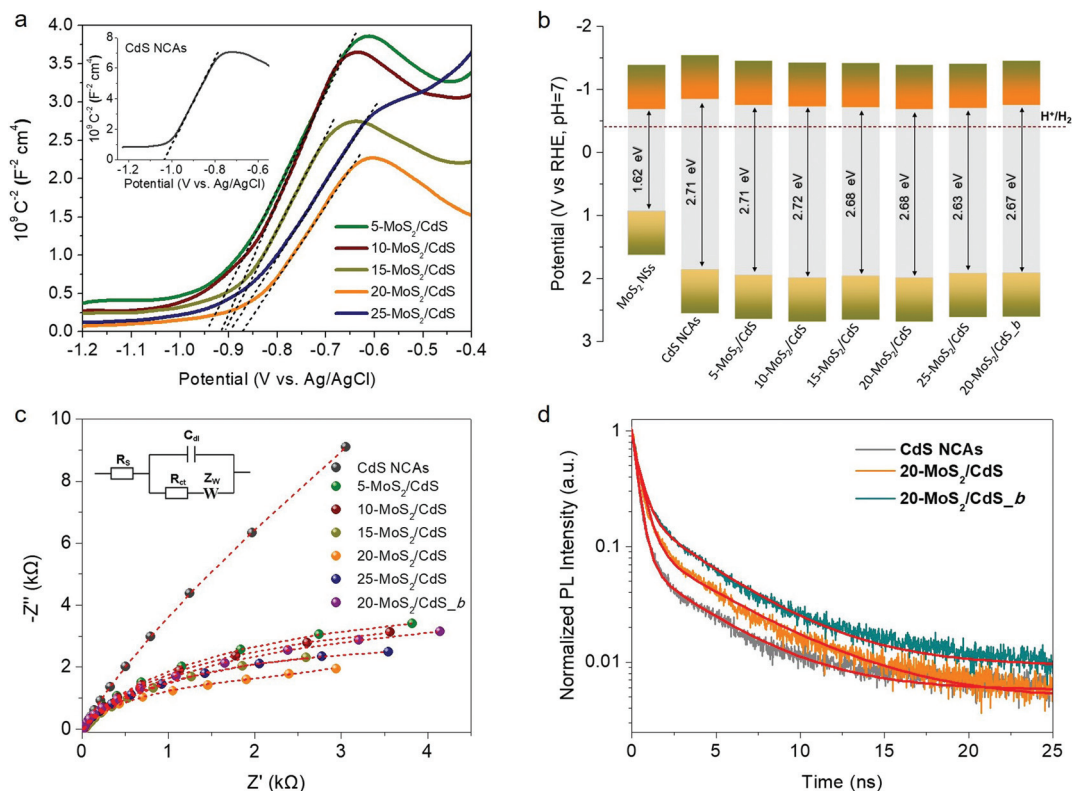
### 3.3 Role of $MoS_2$ NSs in the photochemical activity

To rationalize the role of  $MoS_2$  NSs on the catalytic performance and to better understand the charge-transfer dynamics at the nanoscale  $MoS_2$ /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  $MoS_2$ /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  $MoS_2$ /CdS NCA catalysts range from  $-0.68 \text{ V}$  to  $-0.75 \text{ V}$ , whereas the  $E_{FB}$  position of CdS NCAs locates at  $-0.84 \text{ V}$ ; all

**Table 2** Electrochemical results obtained from EIS analysis for  $MoS_2$  NSs and different mesoporous  $MoS_2$ -modified CdS catalysts

Catalyst	$E_{FB}$ (V vs. RHE, pH 7)	$E_{VB}$	Donor density ( $N_D, \text{cm}^{-3}$ )	$R_{ct}$ (k $\Omega$ )
$MoS_2$ NSs	-0.68	0.94	$6.89 \times 10^{17}$	
CdS NCAs	-0.84	1.87	$5.87 \times 10^{16}$	13.5
5- $MoS_2$ /CdS	-0.75	1.96	$1.17 \times 10^{17}$	6.3
10- $MoS_2$ /CdS	-0.72	2.00	$1.22 \times 10^{17}$	5.6
15- $MoS_2$ /CdS	-0.71	1.97	$1.43 \times 10^{17}$	5.3
20- $MoS_2$ /CdS	-0.68	2.00	$1.70 \times 10^{17}$	2.7
25- $MoS_2$ /CdS	-0.69	1.94	$1.65 \times 10^{17}$	4.5
20- $MoS_2$ /CdS <sub>b</sub>	-0.75	1.92	$1.13 \times 10^{17}$	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  $MoS_2$ -modified CdS NCAs (inset: the Mott–Schottky plot of CdS NCAs). (b) Band-edge positions and (c) Nyquist plots for  $MoS_2$  NSs, CdS NCAs and various  $MoS_2$ -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  $MoS_2$ -modified CdS NCAs with 20 wt%  $MoS_2$  NSs (20- $MoS_2$ /CdS) and  $MoS_2$  bulk flakes (20- $MoS_2$ /CdS<sub>b</sub>).





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  $\text{MoS}_2$  NS content on the surface of CdS. Moreover, as deduced from the magnitude of the slopes of  $1/C_{SC}^2$  vs.  $E$  plots,  $\text{MoS}_2$  modification also increases the donor density ( $N_D$ ) of the catalysts. The measured  $N_D$  values for the  $\text{MoS}_2/\text{CdS}$  NCAs range from  $1.17$  to  $1.70 \times 10^{17} \text{ cm}^{-3}$ , which are significantly larger than that obtained for the unmodified CdS NCAs ( $5.87 \times 10^{16} \text{ cm}^{-3}$ ), as shown in Table 2. The observed anodic shift at the  $E_{FB}$  position of  $\text{MoS}_2/\text{CdS}$  NCAs can be interpreted by the lower  $E_{FB}$  of  $\text{MoS}_2$  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  $\text{MoS}_2/\text{CdS}$  junction (upon contact) that drives electrons from CdS to  $\text{MoS}_2$  until the Fermi levels reach equilibrium; for heavily n-doped semiconductors (such as CdS NCs and  $\text{MoS}_2$  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  $\text{MoS}_2$ . 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  $\text{MoS}_2/\text{CdS}$  junctions, which is in line with the above Mott–Schottky analysis results.

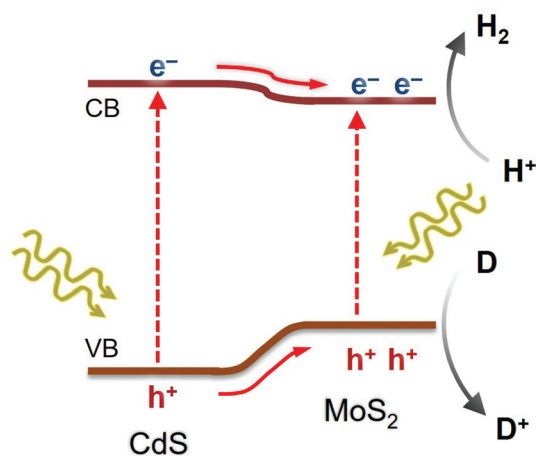
Moreover, the charge transfer properties of the prepared catalysts were also studied by EIS Nyquist plots. Fig. 5c shows the Nyquist plots of pristine and  $\text{MoS}_2$ -modified CdS samples recorded in the  $0.5 \text{ M Na}_2\text{SO}_4$  electrolyte under open circuit conditions over a frequency span from  $1 \text{ Hz}$  to  $1 \text{ MHz}$ . 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  $\text{MoS}_2$ -modified catalysts have  $R_{ct}$  values of  $\sim 2.7$ – $6.3 \text{ k}\Omega$ , which are significantly lower than that of CdS NCAs ( $\sim 13.5 \text{ k}\Omega$ ) (Table 2), suggesting a more favorable charge transfer across the  $\text{MoS}_2/\text{CdS}$  junctions. Subsequently, this process contributes to the improved photocatalytic efficiency of the  $\text{MoS}_2$ -modified catalysts. It is worth mentioning that among diverse  $\text{MoS}_2/\text{CdS}$  catalysts,  $20\text{-MoS}_2/\text{CdS}$  NCAs possessed the lowest resistance in carrier mobility ( $\sim 2.7 \text{ k}\Omega$ ), which associates well with their remarkable photocatalytic  $\text{H}_2$  evolution performance. Notably, the bulk reference catalyst ( $20\text{-MoS}_2/\text{CdS}_b$ ) manifests a relatively higher  $R_{ct}$  value ( $\sim 5.2 \text{ k}\Omega$ ) than the nanostructured analogue ( $20\text{-MoS}_2/\text{CdS}$ ), 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  $20\text{-MoS}_2/\text{CdS}_b$  is lower than that of  $20\text{-MoS}_2/\text{CdS}$  according to the Mott–Schottky equation ( $1.13 \times 10^{17}$  vs.  $1.70 \times 10^{17} \text{ cm}^{-3}$ ), suggesting an inferior electronic contact between the bulk  $\text{MoS}_2$  layers and CdS nanostructure.

This observation agrees with Nyquist charge-transfer resistance measurements and confirms that the nanoscale dimensionality of the  $\text{MoS}_2$  layers has a prominent impact on the charge separation dynamics at the  $\text{MoS}_2/\text{CdS}$  interfaces. This is also elaborated by the photoluminescence (PL) study of catalysts under  $375 \text{ nm}$  excitation wavelength. As shown in Fig. S11,† the  $20\text{-MoS}_2/\text{CdS}$  catalyst shows a weaker PL emission at  $\sim 460 \text{ nm}$  (corresponding to the near band-edge excitonic relaxation) than the pristine CdS and bulk  $20\text{-MoS}_2/\text{CdS}_b$  samples, indicating better mitigation of electron–hole recombination by the nanoscale  $\text{MoS}_2/\text{CdS}$  junctions.

The dynamics of charge-carrier recombination in these materials was also investigated using time-resolved photoluminescence (TRPL) decay measurements upon monochromatic excitation at  $375 \text{ nm}$ . 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  $\alpha_i$  denotes the amplitude fractions ( $\sum_i \alpha_i = 1$ ) and  $\tau_i$  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 ( $\tau_{av}$ ) of mesoporous CdS,  $20\text{-MoS}_2/\text{CdS}$  and  $20\text{-MoS}_2/\text{CdS}_b$  catalysts were determined as  $3.12$ ,  $4.15$  and  $3.46 \text{ ns}$ , respectively; the fitting parameters of the PL decay curves are listed in Table S3.† Once again, TRPL results evidence that the formation of  $\text{MoS}_2/\text{CdS}$  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  $\text{H}_2$  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  $0.18 \text{ V}$ ) in  $\text{MoS}_2$ -modified catalysts compared to the pristine CdS. The observed positive shift in the onset potential with  $\text{MoS}_2$  deposition can be ascribed to the increased charge separation at the  $\text{MoS}_2/\text{CdS}$  interface and the lower energy barrier for  $\text{H}_2$  evolution at the  $\text{MoS}_2$  edge sites. Moreover, in comparison to the bulk reference catalyst ( $20\text{-MoS}_2/\text{CdS}_b$ ),  $20\text{-MoS}_2/\text{CdS}$  NCAs exhibited a quick current response, implying more rapid reaction kinetics caused by the low dimensionality of  $\text{MoS}_2$  layers and increased amount of active  $\text{MoS}_2$  edge-sites. For example, the current density at an applied voltage of  $-1.1 \text{ V}$  versus  $\text{Ag}/\text{AgCl}$  is greatly enhanced from  $0.65 \text{ mA cm}^{-2}$  to  $1.34 \text{ mA cm}^{-2}$  with  $20\text{-MoS}_2/\text{CdS}_b$  and  $20\text{-MoS}_2/\text{CdS}$  catalyst, respectively.

According to the above analysis, a possible reaction mechanism for the photocatalytic  $\text{H}_2$  evolution reaction by  $\text{MoS}_2/\text{CdS}$  nanocatalysts was suggested, as presented in Scheme 1. Light irradiation first excites the electrons from the VB of CdS and  $\text{MoS}_2$  to their respective CB, producing photogenerated carriers. Subsequently, due to the lower CB potential of  $\text{MoS}_2$  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 H<sub>2</sub> production on mesoporous MoS<sub>2</sub>/CdS NCAs under visible light excitation.

active edge sites of MoS<sub>2</sub> NSs, where they efficiently reduce H<sup>+</sup> to H<sub>2</sub>. The smooth charge transfer and abated electron–hole recombination, which are effectively induced by the nanoscale MoS<sub>2</sub>/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 MoS<sub>2</sub> oxidize the sacrificial reagents (such as S<sup>2-</sup> and SO<sub>3</sub><sup>2-</sup>). Evidence of such charge-transfer pathways within the MoS<sub>2</sub>/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 MoS<sub>2</sub> NSs (*ca.* 10–15 nm in lateral size) with different compositions have been successfully synthesized by depositing MoS<sub>2</sub> 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, N<sub>2</sub> 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 MoS<sub>2</sub>/CdS nano-interfaces and promotes high H<sub>2</sub> production efficiency of the photoexcited electrons. Evidence for this was provided by photoluminescence and electrochemical impedance spectroscopy. Owing to these unique characteristics, the MoS<sub>2</sub>/CdS nanojunction network with the optimized MoS<sub>2</sub> content (20 wt%) attains a 6.3× and 1.4× increase in the H<sub>2</sub> production rate compared to bare mesoporous CdS and the bulk MoS<sub>2</sub>/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 MoS<sub>2</sub>-modified CdS catalysts. The recycling test indicated persistent H<sub>2</sub> 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 MoS<sub>2</sub>/CdS nanoscale junctions and support the potential viability of the present MoS<sub>2</sub>-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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