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Mn doped CdS passivated CuInSe₂ quantum dot sensitized solar cells with remarkably enhanced photovoltaic efficiency

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Here we address the optoelectronics properties of CuInSe₂ and Mn doped CdS quantum dot (CISE/Mn-CdS QD) passivated layer solar cells. CISE/Mn-CdS QD solar cells exhibit a remarkably enhanced optical absorption in the visible region as well as monochromatic incident photon-to-electron conversion efficiency (IPCE) (up to 40–90%), which present the interesting photovoltaic efficiency of 3.96% under one sun illumination. Such remarkable optical behavior of the CISE/Mn-CdS QD passivated solar cells is explained by an optical mechanism based on an energy band model and valid physical laws.

1. Introduction

Quantum dot sensitized solar cells (QDSSCs) own many multi-dimensional characteristics, such as size tunable band gaps, ultrafast electron transfer, multi-exciton generation (MEG), high absorption and surpassing the Shockley–Queisser limit (32%), which have attracted the profound attention of researchers over the last few decades.^{1,2} Based on these exciting properties, Cd and Pb-based QDSSCs (CdS, CdSe, CdS/CdTe and PbS, *etc.*) have been found to exhibit a power conversion efficiency (PCE) in the range of 4–9%.^{3–7} Zhong's group has obtained a PCE of 9.73% for a metal hydroxide coated CdSeTe photoanode.⁸ Furthermore, an enhanced PCE of 11% has been reported for the CdSeTe QDSSCs with a carbon counter electrode.⁹ However, the toxic natures of Cd and Pb and the midgap trapping in Pb limit their commercial applications.^{10,11}

In order to solve these problems, researchers have shown their devotion towards the ternary semiconductor quantum dots (TQDs) in photovoltaic devices due to their non-toxic nature, narrow band gap, broad band light harvesting and low cost, such as CuInS₂, CuInSe₂, AgInS₂, *etc.*^{12–14} Among these materials, CuInSe₂ quantum dots (CISE QDs) with the narrow band gap (1.04 eV) have been found to effectively enhance the photovoltaic performance of the QDSSCs due to the large exciton Bohr's radius, which matches well with the solar spectrum.^{15–18} Till now, various techniques have been reported to

synthesize the CuInSe₂ QDs for using in QDSSCs,^{19–21} which include the controllable solvothermal method by incorporating the use of diphenyl phosphine selenide as Se precursor¹⁷ and oleylamine as a solvent and capping ligand.²⁰ Recently, Zhong's group have incorporated Zn in the CISE to obtain the absorption spectra at 1000 nm with PCE at the range of 9%.²² Efficiency has been further enhanced to be 12% by nitrogen doped mesoporous carbon counter electrode in Zn–Cu–In–Se quantum dots sensitized solar cells.²³ However, these QDSSCs still suffer from the issues of low efficiency and poor stability under drastic condition comparing with the conventional solar cells. Therefore, some new areas are required to obtain stable and efficient structure in QDSSCs. These solar cells still need to obtain the optimal structure and fulfill the demanding of the industrial applications.

In order to improve the optical and electrical properties of QDSSCs, various deposition approaches have been employed for the loading and passivation of QDs on the surface of TiO₂ matrix. *In situ* chemical bath deposition (CBD), successive ionic layer adsorption and reaction (SILAR) method,^{24–26} and *ex situ* assembly linking technique⁴ are commonly applied techniques. The surface passivation of QDs by semiconductor materials with large band gap such as CdS and ZnS is carried out to enhance the stability of QDs surface and photovoltaic performance by the prevention of leakage current, which delivers a significant power conversion efficiency (PCE),^{20,27,28} and high open circuit voltage (*V*_{oc}).¹³ The advantage of the Mn²⁺ doping creates an additional electronic state in the CdS host material, which reduces electron–hole recombination and increases a short current density of the QDSSCs. Furthermore, the Mn has dorbital (⁴T₁–⁶A₁) spin and orbital forbidden states, which increase the life time of the excited electron.²⁹ In addition, this type of the engineering structure controls the interfacial charge recombination and updates PCE of the cell. The midgap state

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originated from Mn doping induces the electron to be trapped, which screens them from the recombination with holes and oxidized polysulfide electrolyte.²⁹ Moreover, the addition of the dopant materials elevates the PCE value upto 5.4–9.0% by changing the energy level in case of Mn doped CdS/CdSe and ZnS/SiO₂ QDSSCs.^{27,29} Therefore, the engineering of Mn-doped material will greatly enhance the PCE of the QDSSCs.

Herein, we employ a low temperature and simple thermolysis method, followed by the capping of bifunctional molecule to prepare mercaptopropionic acid (MPA) capped CISE QDs, and the CISE QDs are sensitized on TiO₂ matrix by *in situ* linker assisted technique. Furthermore, the passivation layer of Mn-doped CdS (Mn-CdS) QDs over CISE QDs/TiO₂ matrix is carried out by SILAR method.²⁷ The doping produces a stable structure under severe temperature condition and fast electron transfer. These CISE/Mn-CdS QD passivated QDSSCs exhibit significant optical and electrical properties under one sun illumination.

2. Experimental

2.1 Synthesis of CuInSe₂ QDs

All chemicals were analytically grade and were used without further purifications. CuInSe₂ QDs were synthesized by a modified thermolysis method as reported in literature.¹⁸ Briefly, 10 mL oleylamine (OLA), 1 mmol CuI (0.190 g) and 1 mmol InCl₃ (0.220 g) were simultaneously added to a three neck flask and degassed for 30 minutes. Then, the temperature was increased to 180 °C under the argon flow and 2 mmol DPP/Se (1 : 1 ratio) was injected and the temperature was maintained at 180 °C for a further 10 minutes. The mixed solution was cooled to room temperature and washed by centrifugation through methanol and hexane for three times. The precipitates were dried at 100 °C in vacuum oven to obtain the CISE QDs powder.

The as-prepared CISE QDs were dispersed in fresh hexane for sensitization. To obtain MPA capped QDs, CISE QDs/methanol solution (1 mL) was dissolved in methanol solution of MPA and NaOH, which was stirred for 30 minutes, then deionized water was added to the mixture and maintained stirring for further 20 minutes. The precipitates were finally washed through centrifugation in excess acetone and deionized water to obtain water soluble MPA capped CISE QDs.

2.2 Device fabrication

TiO₂ nanoparticle working electrodes were prepared by screen printing technique as reported in literature.³⁰ Briefly, fluorine doped tin oxide (FTO) glass was first treated with TiCl₄ solution at 70 °C for 3 minutes. Back scattering layer was prepared from P25 paste, followed by the top layer of 400 nm TiO₂ nanoparticles. Finally, the TiO₂ thin films were sintered at 450 °C in the muffle furnace, and then dipped in CISE QDs solution for 24 hours to complete the sensitization. These CISE QDs sensitized TiO₂ films were washed by water and dried under vacuum.

MPA capped CISE QDs were sensitized on TiO₂ films followed by the passivation of Mn-CdS layer by successive ionic layer

adsorption and reaction (SILAR) method. In short, 0.1 M cadmium nitrate in methanol was used as cation source and 0.1 M sodium sulfide in 1 : 1 methanol and water as anion source. To incorporate Mn doping, manganese acetate (0.075 M) was mixed with cadmium nitrate. The CISE QDs sensitized films were immersed for one minute in Mn doped Cd precursor followed by rinsing with methanol, and then immersed in S precursor followed by rinsing with methanol, which counts as one SILAR cycle and repeats the process for 3 cycles. Finally the devices were dried under Ar atmosphere to remove the organic solvents.

The QDs co-sensitized solar cells were fabricated by assembling the sensitized TiO₂ photoanode and brass based Cu₂S counter electrode with a binder clip, separated by a 60 mm Scotch tape spacer. Polysulfide electrolyte solution with composition of Na₂S (2.0 M), S (2.0 M) powder and KCl (0.2 M) was injected into the cells device.

2.3 Characterization and measurements

Structure and morphology were measured by X-rays diffraction analysis (XRD, Pertpro, PANalytical, Netherlands) using Cu K α , field emission scanning electron microscope (FESEM, JSEM-5610LV, Japan) and the high resolution transmission electron microscope (HRTEM, JEM-2100F, JEOL, Japan), respectively. The optical characterization of the samples was carried out by using UV-vis absorption spectra (UV-2550, Shimadzu, Japan). The photocurrent density–voltage (*J*–*V*) curves of the QDSSCs were measured by the Keithley 4200 semiconductor characterization system (Keithley Instruments, USA) under simulated AM 1.5 illumination (Newport 91160, 300 W xenon lamp, USA). The incident photon-to-current conversion efficiency (IPCE) was measured using Newport's IPCE Measurement Kit under the monochromatic light from a monochromator (Newport). The electrochemical impedance spectra (EIS) were measured by using an impedance analyzer (Auto lab PGSTAT320N) at open-circuit potentials under AM 1.5 illumination.

3. Results and discussions

XRD pattern of CISE QDs is shown in Fig. 1(a), and the diffraction peaks are indexed to the chalcopyrite (tetragonal) CuInSe₂ QDs (JCPDS 00-051-1221) with all major peaks reflected from (112), (204), (312) and minor peak of (211) crystal plane at angle 2θ of 26.77°, 44.43°, 54.64° and 31.01°, respectively, which is the stable phase of CISE QDs at room temperature. The significant broadening in the peaks represents the small size of CuInSe₂ QDs. Applying Scherrer's formula, the average size of QDs is calculated to be 5 nm.³¹ Fig. 1(b) and (c) illustrate the top and cross sectional views of the as-prepared film with the thickness of 6 μ m. Average size of as-synthesized QDs is 5 nm as shown in the Fig. 1(d), which is in good agreement with the XRD analysis. The HRTEM image displays high crystallinity of the as-synthesized CISE QDs in Fig. 1(e), and the inter-planar spacing is about 0.362 nm, which corresponds to the (112) crystal plane of the tetragonal structure.



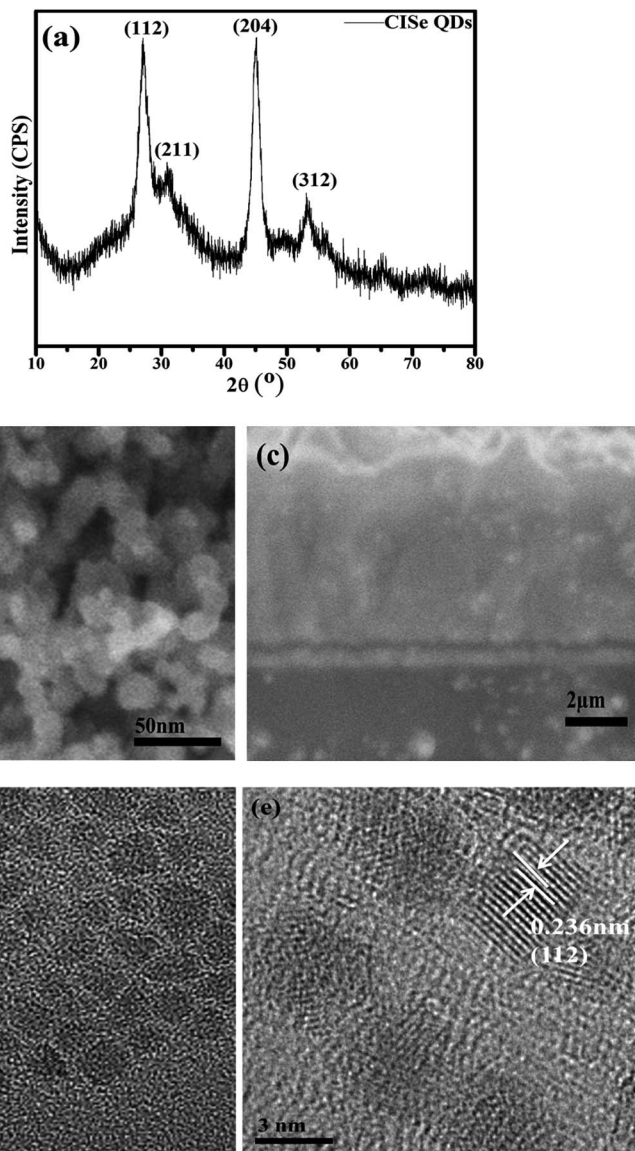


Fig. 1 Characteristic of the as-prepared CuInSe_2 quantum dots: (a) XRD pattern, (b) top and (c) cross-sectional views of FESEM images, (d) TEM image, (e) HRTEM image.

UV-vis spectra of the TiO_2 photoanodes sensitized by CISE, CdS, Mn-CdS and CISE/Mn-CdS QDs are shown in Fig. 2(a). After the sensitization of CISE QDs, the absorption spectrum of the TiO_2 films is significantly enhanced both in ultraviolet and visible light regions. The obvious blue shift is observed after the sensitization of CISE QDs with the main peak centered at 450 nm, while the absorption onsets of the TiO_2/CdS and $\text{TiO}_2/\text{Mn-CdS}$ occur approximately at 560 and 580 nm, respectively, which show the characteristics spectra toward red shift compared with that of the bulk CdS material. The CISE/Mn-CdS QD passivated electrode exhibits red-shift absorption at the wavelength range of 650–700 nm. Mn-CdS passivation layer has an absorption onset at 580 nm, the red-shift phenomenon must result from the light absorption of the CISE QDs. The passivation enhances the absorption spectra towards long wavelength region comparing with those of pure TiO_2 and CISE/ TiO_2 matrix.

This red-shift may indicate that the QDs charge carriers wave functions tunnel into the surrounding CdS shell. The broad spectrum reveals the heterogeneity of the CISE and Mn-CdS QDs sizes.³² Furthermore, UV-vis spectra are also measured to reveal the band gaps of the as-prepared samples. Herein, using Tauc relation (eqn (1)), the optical band gap of CISE QDs is calculated as 1.38 eV and other corresponding materials are shown in Fig. 2(b).^{33,34}

$$\alpha h\nu = A(h\nu - E_g)^{\frac{n}{2}} \quad (1)$$

where α is absorption coefficient and given by $\alpha = 2.303$, and $A = \log(T/d)$ (d is the thickness of the sample and T is the transmission), $h\nu$ is the photon energy. Further, the bands gaps of TiO_2 , CdS, Mn-CdS and CISE/Mn-CdS are 3.1, 2.3, 2.2 and 1.6 eV, respectively.



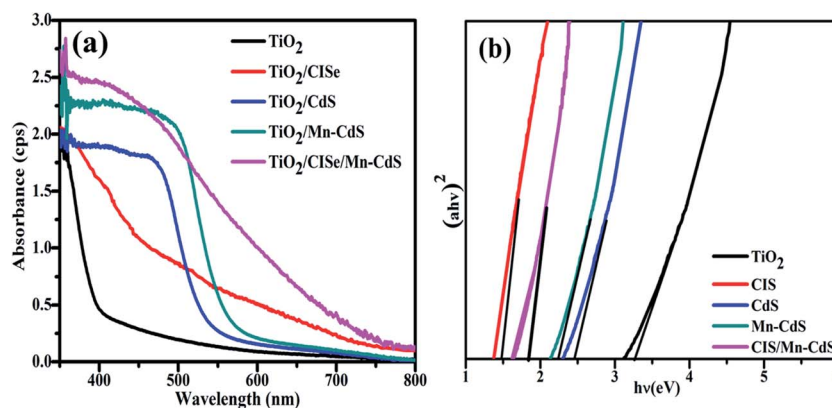


Fig. 2 (a) UV-vis spectra of TiO_2 , CISe, CdS, Mn-CdS and CISe/Mn-CdS QDs sensitized solar cells, (b) the Tauc plots for the determination the band gap.

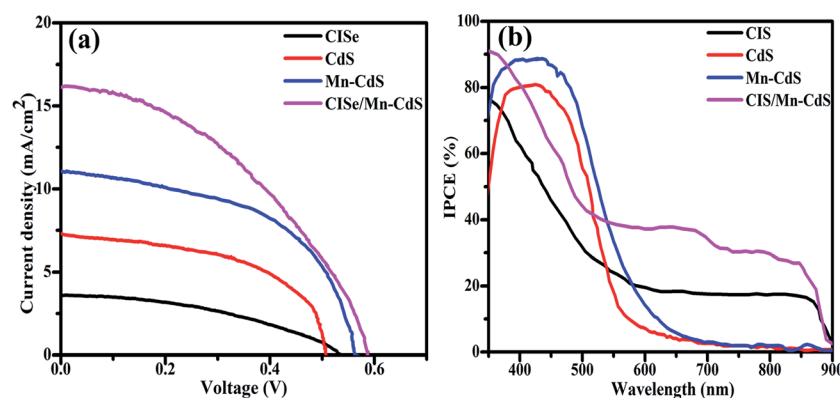


Fig. 3 The photovoltaic performance of the CISe, CdS, Mn-CdS and CISe/Mn-CdS sensitized solar cells under AM 1.5 sunlight: (a) J - V curves, (b) IPCE curves.

The photovoltaic performance is determined from the current-voltage (J - V) characteristics of the assembled devices as shown in the Fig. 3(a). The photovoltaic parameters of these solar cells, such as the open circuit potential (V_{oc}), short circuit current density (J_{sc}), fill factor (FF) and power conversion efficiency (PCE), are listed in Table 1. The photovoltaic efficiencies of CISe QDs, CdS QDs, Mn-CdS QDs and CISe/Mn-CdS QD passivate solar cells are 0.80, 1.98, 3.08 and 3.96%, respectively. For CISe/Mn-CdS QD passivate solar cells, the decreasing of the FF to 41% clearly indicates the reduction in the internal charge recombination and defect state in the QDSSCs.¹⁸ Moreover, the increasing in the current density reveals the high charge transportation in photovoltaic device. CISe/Mn-CdS QD passivate solar cells exhibit an enhanced open circuit voltage of 0.59 V, which substantiates a high charge extraction in the alloy structure. However, the obtained photovoltaic efficiency in the present work is still lower comparing with some of similar works.^{8,18,22,29}

Normally, the PCE value of QDSSCs is greatly influenced by the light absorption ability, which is related with the sensitization ratio of QDs as well as the size and morphology of the photoanode film. In our case, as the passivation of Mn-CdS QDs inside the pores of CISe QDs influences the electrolyte diffusion

inside the photoanodes, which leads to lower the charge diffusion and ultimately limit the cell performance.³⁵ The lower diffusion length causes the decreasing of the number of the extracted electrons to the TiO_2 matrix. Conduction band level of CISe QDs is lower than that of n-type Mn-CdS QDs, which reduces the charge transfer across the CISe/Mn-CdS interface to cause a low photovoltaic performance.³⁶ In our case, the SILAR and CBD deposition may lead to the undesired defect state due to the broad size distribution, thereby, the undesired surface defects are created, which leads to the trapping of charge carrier and the decreasing of the PCE value.^{29,37}

IPCE is measured for electrons extraction/collection excited by incident photons as shown in Fig. 3(b). IPCE values of CISe, CdS, Mn-CdS and CISe/Mn-CdS QDSSCs show high values even in longer wavelength regions, and good consistency with the absorption spectra as shown in Fig. 2(a), which confirms the large number of electrons transport to the TiO_2 . The lower IPCE values of the CdS and Mn-CdS sensitized solar cells are due to their weak response at low energy wavelength.³⁸ The enhancement in the IPCE value is noted for CISe/Mn-CdS QDSSCs with a maximum IPCE in the range of 40–90% with the obtained power conversion efficiency of 3.96%. This high response of incident photon to excite electrons conversion is due to the



Table 1 Photovoltaic parameters of various quantum dots sensitized solar cells extracted from J - V measurements

Type	V_{oc} (V)	J_{sc} (mA cm ⁻²)	FF	Efficiency (%)
CISE QDs	0.53	3.50	43	0.80
CdS QDs	0.51	7.20	52	1.98
Mn-CdS QDs	0.56	10.26	49	3.08
CISE/Mn-CdS QDs	0.59	16.10	41	3.96

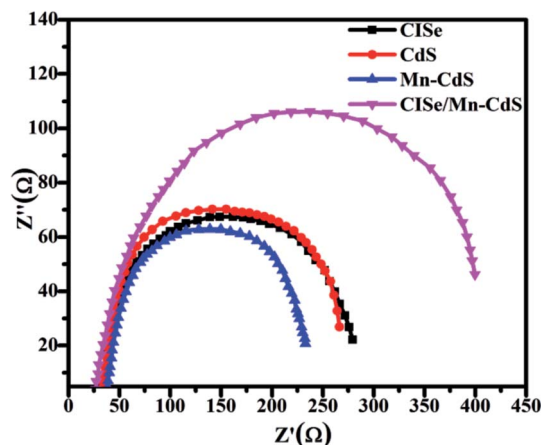
narrow band gap of CISE QDs, which can deduce a high electron injection upon the incident light photon. Moreover, the passivation of Mn-doped CdS QDs improves the performance of electron collection/injection, as the large number of the photo-induced electrons in CISE/Mn-CdS QDSSCs are injected into and then collected efficiently on TiO₂ matrix. The high photo-responses in short and long wavelength regions are due to the combination of narrow and wide band gap CISE/Mn-CdS heterostructure, which absorbs the light in both of ultraviolet and visible regions. The photoresponse can be calculated by the following equation.³⁹

$$\text{IPCE} = \text{LHE} \times \phi_{\text{inj}} \times \eta_{\text{cc}} \quad (2)$$

where LHE is the light harvesting efficiency, ϕ_{inj} is the electron injection efficiency and η_{cc} represents the charge collection efficiency.³⁹ Superior light harvesting and photoelectrochemical performance of CISE/Mn-CdS QDSSCs is another breakthrough for achieving the high photovoltaic efficiency.

QD loading ratio on the TiO₂ surface is an important factor for high photovoltaic performance. The advantage of *in situ* chemicals bath deposition method is the insertion of the small size QDs in the space between TiO₂ nanoparticles, thereby reduces the defect levels and electron-hole recombination, which ultimately enhances the photovoltaic performance. Firstly, MPA is a short chain ligand, which acts as the linker molecule between QDs and TiO₂ thin film through a strong interaction of (Ti-O-Ti) with carboxyl group (COOH). Secondly, SH group of MPA firmly interacts with Cu ion through S-metal junction.⁴ Thirdly, according to soft and hard acid basis theory, Se²⁻ ions have a strong binding with soft ions-SH from MPA.⁴⁰ The MPA capped and water soluble CISE QDs have a strong affinity to the TiO₂ molecules. Moreover, under the simulated sunlight irradiation, the MPA linking effectively reduces the blocking of photo-induced electrons transfer and the recombination possibility of photo-induced electrons and holes, which results in the improvement of photovoltaic conversion efficiency.⁴¹

Electrical impedance spectra (EIS) expose the dynamics of bound or mobile charge in bulk or interfacial region inside the solar cells. The EIS measurements are carried out under the dark condition at different forward biases ranging from 0 to -0.5 V. Nyquist plots of the cells at forward biased -0.5 V are shown in the Fig. 4. Comparing with the Nyquist plots of CISE, CdS and Mn-CdS QDSSCs, the CISE/Mn-CdS QDSSCs reveal a high R_{rec} value, suggesting a considerably larger electron transport resistance between the photo-electrode and polysulfide electrolyte.³² The R_s

**Fig. 4** EIS plots of CISE, CdS, Mn-CdS and CISE/Mn-CdS sensitized solar cells.

values of all the electrodes are nearly same, which is attributed to the fact that the passivation has no effect on the interfaces. Small semicircle of Mn-CdS QDSSCs indicates the favor of electron transfer from Mn-CdS QDs to the TiO₂ matrix. The low resistance feature of the Mn-CdS QDs may be due to the forbidden d-d orbital transition, which impules the transportation electron and lowers the resistance. The large semicircle of CISE/Mn-CdS QDSSCs shows the high interface resistance, which is the sum of the resistances among Mn-CdS QDs, CISE QDs and TiO₂ interfaces in the QDSSCs and increases with the increasing of the film thickness. However, the surface defect and dangling bonds of the TiO₂ film are removed by the Mn-CdS growth, which ultimately suppresses the photo-induced electron-hole recombination. In comparison with the other electron deterring approaches, the CISE/Mn-CdS alloy structure in our work exhibits a compelling performance in reducing the charge recombination and improving power conversion efficiency. The high recombination resistance of the Mn-CdS passivated CISE QDSSCs is possibly originated due to the surface trap states, which favor for the recombination to lower the PCE value of the device. The simulated results of the as-prepared samples are listed in the Table 2. These results confirm that the passivation of Mn-CdS QDs on CISE QDSSCs is an effective tool to suppress the charge recombination. It eventually contributes to the excellent properties of QDSSCs.

Under the sun light irradiation with sufficient energy, the photo-induced electrons jumps from valence band (VB) to conduction band (CB) of Mn-CdS QDs. Due to the large band gap offset, the photo-induced electrons transfer from the Mn-

Table 2 Simulated values of the resistance and capacitance of CISE, CdS, Mn-CdS and CISE/Mn-CdS QDSSCs

Sample	R_s (Ω cm ²)	C_μ (mF cm ⁻²)	R_{rec} (Ω cm ²)
CISE QDSSCs	30.54	0.51	285.9
CdS QDSSCs	30.20	0.46	271.4
Mn-CdS QDSSCs	36.30	0.54	223.8
CISE/Mn-CdS QDSSCs	24.80	0.72	443.0



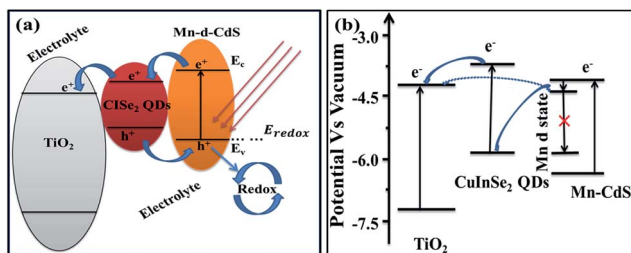


Fig. 5 Schematic diagrams of CISE/Mn-CdS sensitized solar cells: (a) band gap, (b) electron transport path.

CdS CB to CuInSe₂ CB and then to TiO₂ CB, which increases the photocurrent density. Furthermore, the photo-generated electrons from the CB of CuInSe₂ QDs and Mn-CdS QDs with the high absorption of sun light transfer to TiO₂ CB, which corporately increase the current density of the optoelectronic device. Hereafter, it is clearly observed that Mn doping causes the delay of the photo-induced electrons in the excited state, thereby enhances V_{oc} . Schematic diagram of the photo-induced electron flow is enlightened in Fig. 5. Mn doped CdS QDs create some states in the mid-bandgap region due to the d atomic orbitals of Mn (⁴T₁–⁶A₁). However, the transitions between these two energy levels (⁴T₁–⁶A₁) are both spin and orbital forbidden.²⁷ Thereby, the photo-induced electrons stay for a long time at Mn d-states and then are injected to CISE QDs as well as to TiO₂. There are some additional deep electron trap states, which reduce the concentration of mobile carriers in the n-type CdS layer. The generation of the excitons in the CdS QDs causes a compensation of deep trap by ‘photo-doping’,^{42,43} which increases the number of mobile carriers, hence reduces the barrier towards electron transport across the CdS QDs and leads to an increased junction conductance.⁴⁴ Mn-CdS QDs have a higher mobility of photo-induced electrons than CdS QDs, which increases the current density of the Mn-CdS QDSSCs. Furthermore, the long life time of d–d transition in Mn delays the electron–hole recombination in the QDs, which alternately increases the open circuit potential.

4. Conclusion

Water soluble CISE QDs of 5 nm are prepared by a simple thermolysis method. Co-sensitization of CISE/Mn-CdS QDs clearly enhances the optical and electrochemical properties of the solar cells. The excellent architecture of CISE/Mn-CdS demonstrates the outstanding light-harvesting behavior of the QDSSCs. MPA-capping improves the QDs loading on TiO₂ photoanode. The combination of *in situ* linker and SILAR method in the fabrication of CISE/Mn-CdS alloy architecture suppresses the photo-induced electron–hole recombination; as a result, the photovoltaic efficiency of 3.96% is achieved for CISE/Mn-CdS QDSSCs.

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