

Electronic Supporting Information (ESI)

Highly Efficient Colloidal $\text{Mn}_x\text{Cd}_{1-x}\text{S}$ Solid Solution Nanorods for Photocatalytic Hydrogen Generation

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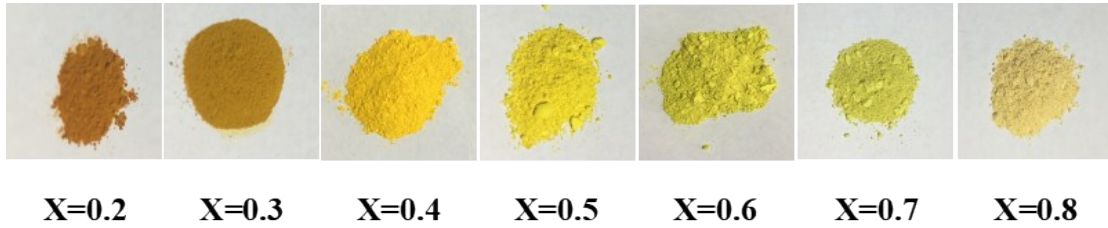


Fig. S1 Digital photographs of synthesized $Mn_xCd_{1-x}S$ samples.

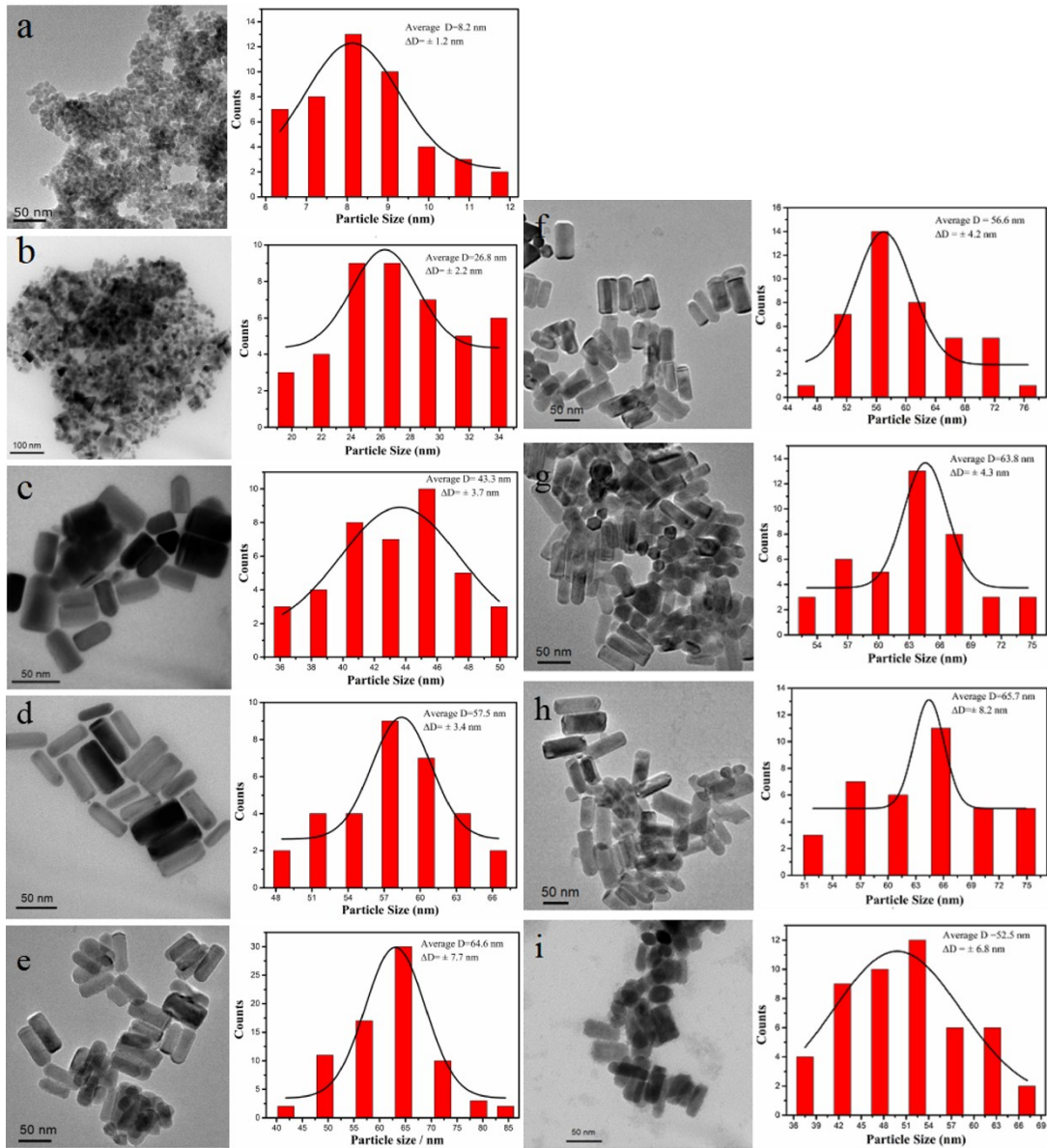


Fig. S2 TEM and Size distribution histogram of $Mn_xCd_{1-x}S$ solid solutions. (a. $x=0$; b. $x=0.2$; c. $x=0.3$; d. $x=0.4$; e. $x=0.5$; f. $x=0.6$; g. $x=0.7$; h. $x=0.8$; i. $x=1.0$)

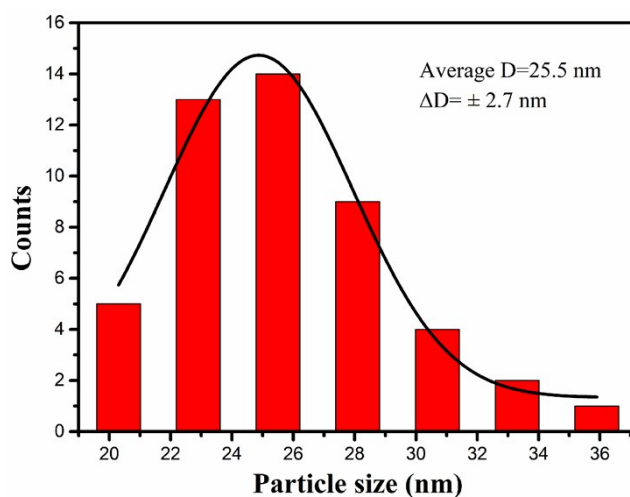


Fig. S3 Size distribution histogram of $Mn_{0.5}Cd_{0.5}S$ nanorods (top of view).

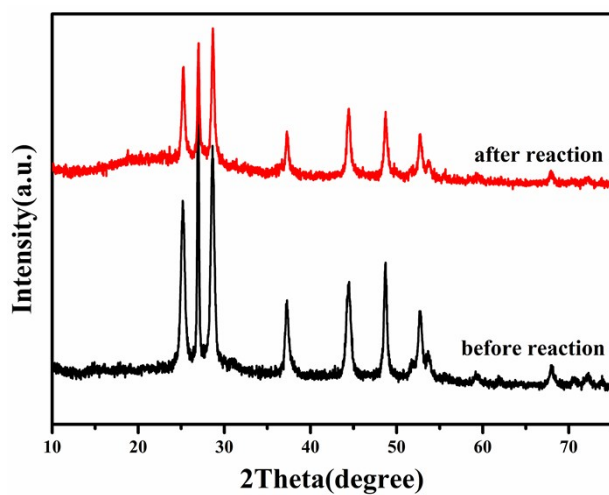


Fig. S4 XRD patterns of the $Mn_{0.5}Cd_{0.5}S$ solid solution photocatalyst before and after the photocatalytic reactions.

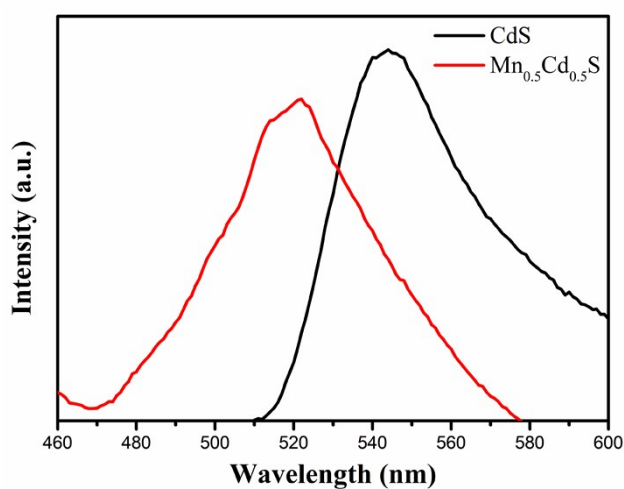


Fig. S5 Photoluminescence spectra of CdS (black) and $Mn_{0.5}Cd_{0.5}S$ (red) in solution.

Table. S1 Comparison of H₂ evolution rate with other Mn-Cd-S solid solutions.

No.	Photocatalyst	Rate of H ₂ Evolution	Method	Reference
1	Mn _{0.2} Cd _{0.8} S/CoP ₃	29530 μmolg ⁻¹ h ⁻¹	Hydrothermal method	1
2	Mn _x Cd _{1-x} S	355 μmolg ⁻¹ h ⁻¹		2
3	Mn _x Cd _{1-x} S	10900 μmolg ⁻¹ h ⁻¹		3
4	Mn _{0.25} Cd _{0.75} S/MoS ₂	12470 μmolh ⁻¹ g ⁻¹		4
5	Mn _{0.8} Cd _{0.2} S/g-C ₃ N ₄	4000 μmolg ⁻¹ h ⁻¹		5
6	Mn _x Cd _{1-x} S/NiS	8386 μmolh ⁻¹ g ⁻¹		6
7	Cu _{2-x} S/Mn _{0.5} Cd _{0.5} S/MoS ₂	13752 μmolg ⁻¹ h ⁻¹		7
8	Ni doped Mn-Cd-S	1020 μmolg ⁻¹ h ⁻¹		8
9	Ag doped Mn-Cd-S	4400 μmolg ⁻¹ h ⁻¹		9
10	Zn _{1-x} Cd _x S solid solutions	7420 μmolg ⁻¹ h ⁻¹	thermolysis method	10
11	ZnCdS dodecahedral cages	5680 μmolg ⁻¹ h ⁻¹	Sulfurization and ion exchange	11
12	NiS/ Zn _x Cd _{1-x} S/RGO	7514 μmolg ⁻¹ h ⁻¹	Hydrothermal method	12
13	Cu ²⁺ doped In _{2x} Zn _{3(1-x)} S ₃	790 μmolg ⁻¹ h ⁻¹		13
14	MnS/In ₂ S ₃	8360 μmolh ⁻¹ g ⁻¹ (from H ₂ S)	Solvothermal method	14
15	Mn_xCd_{1-x}S	26000 μmolg⁻¹h⁻¹	Colloidal method	This study

Table. S2 PL lifetime fitting results for CdS and Mn_{0.5}Cd_{0.5}S

sample	T1 / ns	T2 / ns
CdS	6.31	123.44
Mn_{0.5}Cd_{0.5}S	7.60	162.38

The PL kinetic curves were fitted with bi-exponential decay function, thus providing two lifetimes, one fast decay component and one slow decay component. Fast decay components were usually associated with non-radiative recombination, while slow decay components were due to the radiative lifetime of free excitons.¹⁵

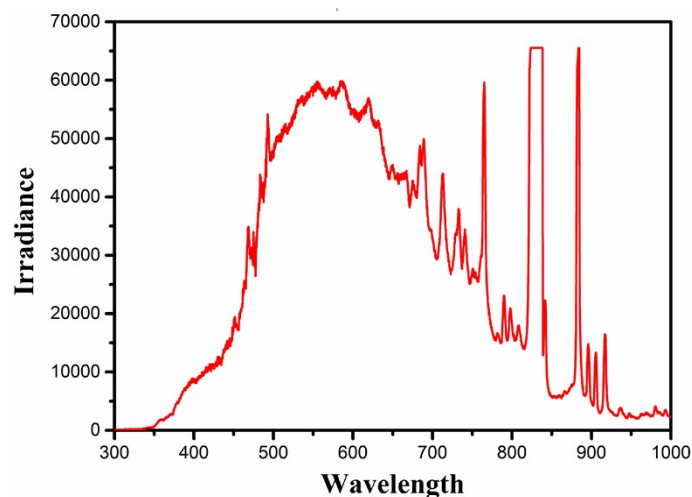


Fig. S6 Emission spectra of Xenon light source. (PLS-SXE300/300UV; PerfectLight, China.)

Reference:

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