

A novel heterogeneous CdS nanoparticles/NiTiO₃ nanorod with enhanced visible-light-driven photocatalytic activity

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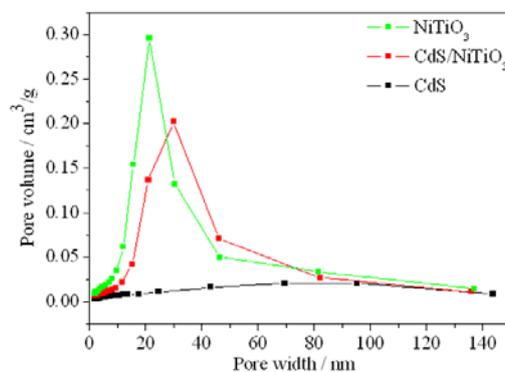


Figure S1. The average pores distribution of NiTiO₃ nanorods, CdS nanoparticles and heterogeneous CdS nanoparticles/NiTiO₃ nanorods.

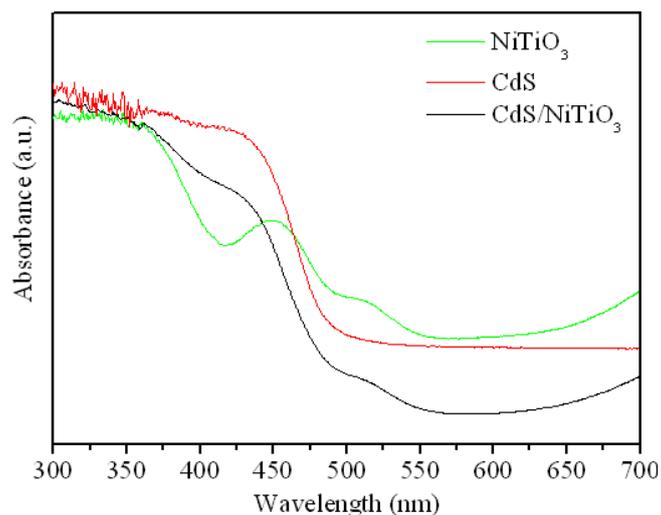


Figure S2. UV-vis diffuses reflection spectroscopy of NiTiO₃ nanorods, CdS nanoparticles and heterogeneous CdS nanoparticles/NiTiO₃ nanorods.

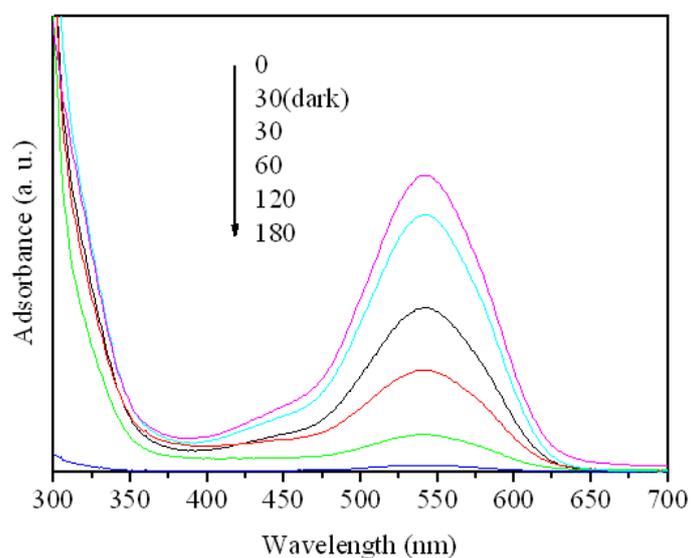


Figure S3. UV-vis spectra of aqueous solutions of Cr (VI) after photocatalytic reaction by the diphenylcarbazide method of heterogeneous CdS nanoparticles/NiTiO₃ nanorods for different times irradiation.

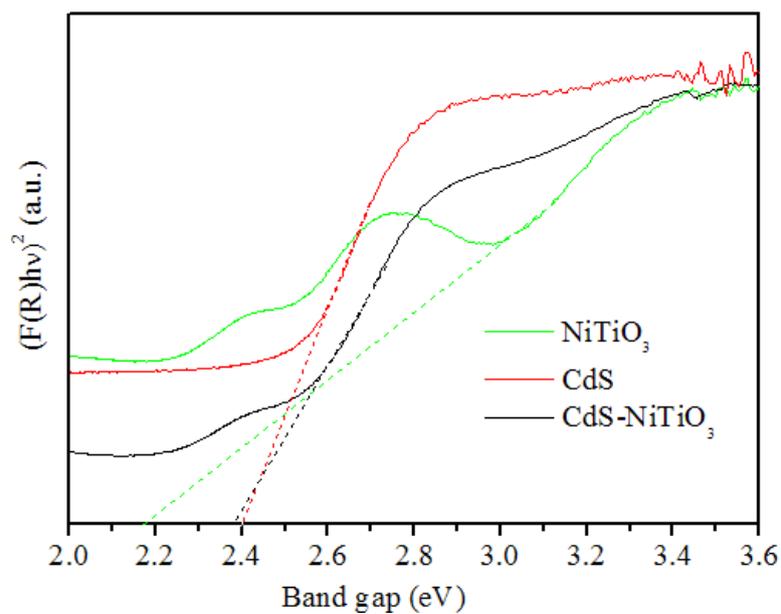


Figure S4. Plots of $(F(R)hv)^2$ versus (hv) for obtaining the band gaps of NiTiO₃ nanorods, CdS nanoparticles and heterogeneous CdS nanoparticles/NiTiO₃ nanorods.

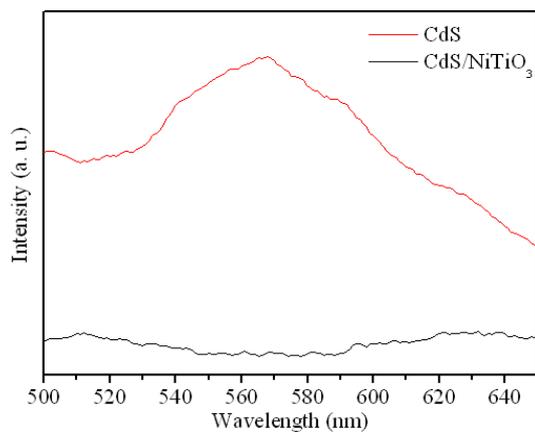


Figure S5. PL spectra of CdS nanoparticles and heterogeneous CdS nanoparticles/NiTiO₃ nanorods.

Table S1. The band gap and conductor band data with respect to Absolute Vacuum Scale (AVS) for common semiconductors.

Semiconductors	CdS	NiTiO ₃	TiO ₂	ZnO	SrTiO ₃	KTaO ₃
Band gap (eV)	2.4	2.18	3.2	3.2	3.4	3.5
Conductor band (eV)	-3.98	-4.70	-4.21	-4.19	-3.24	-3.57

The band structure data was obtained from Ref. 17.

Table S2. The Concentration of Cd²⁺ in the solution after 1h irradiation of photocatalytic reduction of Cr (VI).

Reaction	Photocatalytic reduction of Cr(VI)	
Samples	CdS/NiTiO ₃	CdS
Concentration of Cd ²⁺ in the solution after 1h irradiation of photocatalysis	0.063 mg/L	1.134 mg/L

It was measured by atomic absorption spectrometry (AAS, Thermo Elemental SOLAAR-M, limit of identification: 5 µg L⁻¹). The samples were centrifuged and filtered through 0.22 µm nominal pore-size membrane filters prior to analysis.