

Supporting Information

1. Materials and instruments

1.1 Materials

$\text{Y}(\text{CH}_3\text{COOH})_3 \cdot 4\text{H}_2\text{O}$, $\text{Yb}(\text{CH}_3\text{COOH})_3 \cdot 4\text{H}_2\text{O}$, $\text{Tm}(\text{CH}_3\text{COOH})_3 \cdot 4\text{H}_2\text{O}$, $\text{Gd}(\text{CH}_3\text{COOH})_3 \cdot 4\text{H}_2\text{O}$, oleic acid, 1-octadecene, and polyvinylpyrrolidone (Mw=40,000) were purchased from Sigma-Aldrich. 1,4-benzoquinone, ethylenediaminetetraacetate, and 2-aminoterephthalic acid, and Rhodamine B were purchased from Adamas. NaOH, NH_4F , Chromium nitrate nonahydrate and isopropanol were purchased from Greagent. All the chemical reagents were used as received without further purification.

1.2 Experimental and characterization instruments

The microstructure of NYT, NYTG(3%), and NYTG(3%)/NMC were studied by a transmission electron microscope (TEM, JEM-2100, operated at 200 kV). The crystal structures of all the samples were analyzed by the X-ray diffraction instrument (XRD, Bruker D8 Advance) with a Cu $\text{K}\alpha$ radiation source ($\lambda = 0.15418$ nm) operated at 40 kV and 40 mA. The UV-Vis-NIR diffuse reflectance spectra (DRS) of the as-prepared solid samples were investigated by using a PE Lambda 950 spectrophotometer, which equipped with an integrating sphere. The upconversion luminescence spectra were obtained by using a Fluorescence spectra were recorded on FluoroMax+ spectrometer with an external 980 nm NIR laser. The XPS spectra were performed by Thermo Scientific K- α + XPS spectrometer with the monochromatized Al-K α X-ray source ($h\nu = 1486.8$ eV). The 400 μm X-ray spot was used for XPS analysis. The absorbance of RhB in aqueous solutions was measured using a UH4150 spectrophotometer. Photocatalytic efficiency of RhB decomposition was tested under UV-Visible (<760 nm), near-infrared light (>760 nm) and full-spectra-light

irradiation with a 300 W Xenon lamp (Perfect Light PLS-SXE-300+) that equipped with a cooling system for temperature control. The simulated solar light photocatalysis reaction was carried out using a solar simulator (AM 1.5 G).

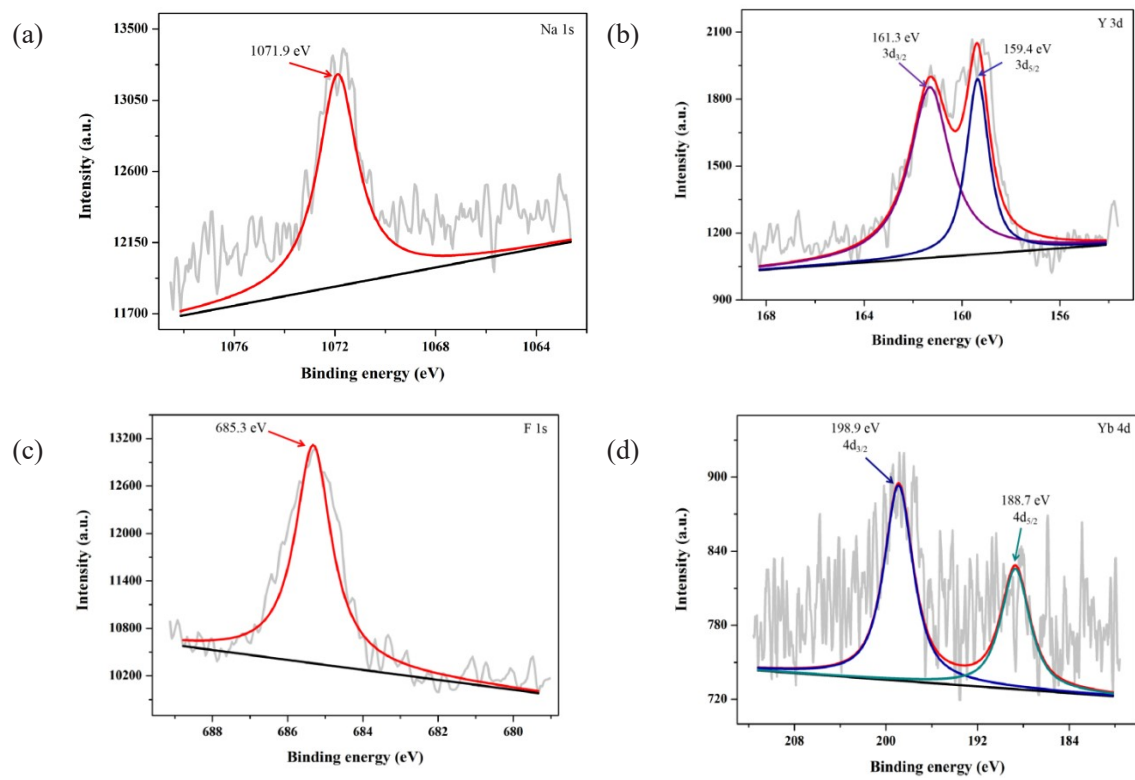


Fig. S1. XPS spectra of NYTG(3%)/NMC: (a) Na 1s, (b) Y 3d, (c) F 1s, and (d) Yb 4d

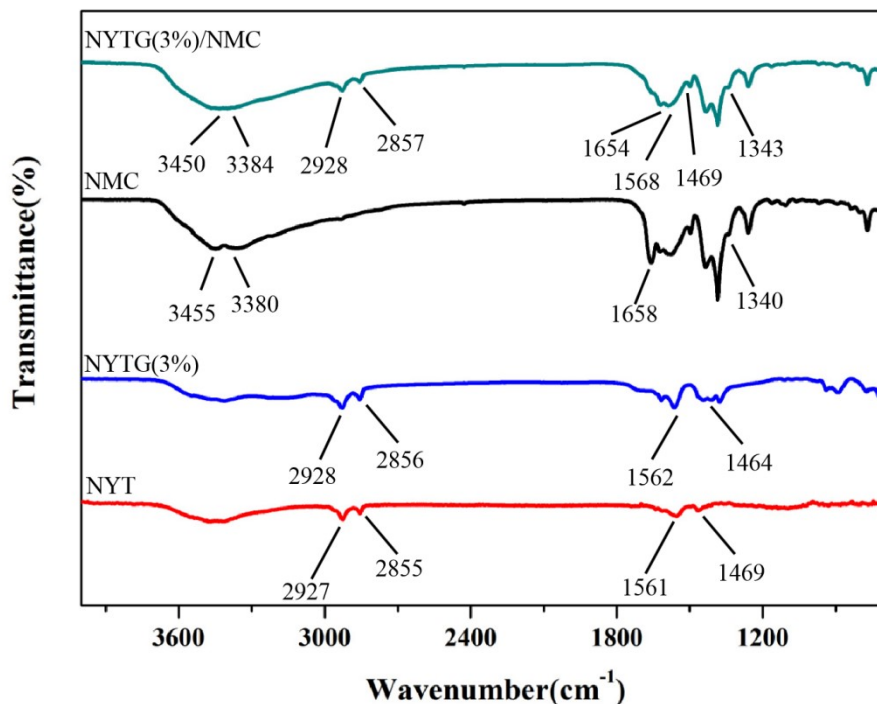


Fig. S2. Fourier transform infrared (FTIR) spectra of NYT, NYTG(3%), NMC, and NYTG(3%)/NMC samples.

Fig. S2 demonstrated the FT-IR spectra of NYT, NYTG(3%), NMC, and NYTG(3%)/NMC samples. For the NYTG(3%)/NMC sample, the peaks of NYTG(3%) and NMC are all observed indicating that NYTG(3%)/NMC was successfully synthesized. The peak at 1658 cm^{-1} was attributed to the N-H bending vibration of the amino groups in NMC. And the tiny peak at 1340 cm^{-1} corresponds to the C-N stretching on benzene ring. Moreover, the double peaks at 3455 cm^{-1} and 3380 cm^{-1} can be assigned to asymmetric and symmetric vibration of the amino groups in NMC. The peaks at around 2928 and 2855 cm^{-1} corresponding to $-\text{CH}_2-$ group and peaks near 1562 and 1464 cm^{-1} corresponding to $-\text{COOH}$ group were clearly observed suggesting that the successful involvement of NYT and NYTG(3%). The introduction of Gd^{3+} ions does not change the infrared peak of upconversion nanoparticles.

Table S1 Photodegradation performance of different upconversion nanoparticles based photocatalysts.

Ln-UCNPs	Coating materials	Light source	Pollutant	Degradation ratio/time (min)	Ref.
NaYF ₄ :Yb,Tm	TiO ₂ , Fe ₃ O ₄	simulated sunlight	MB	75%/50	1
NaYF ₄ :Gd,Si	TiO ₂	simulated sunlight	MB	70%/240	2
BiOBr:Yb,Er 3D Hierarchical Architectures	/	simulated sunlight	RhB	100%/40	3
NaLuF ₄ :Gd,Yb ,Tm@NaLuF ₄ : Gd,Yb	SiO ₂ , Ag, g- C ₃ N ₄	simulated sunlight	RhB	100%/150	4
NaYF ₄ :Yb,Tm @NaYF ₄	Zn _{0.5} Cd _{0.5} S	simulated sunlight	Cr(VI)	98.7%/30	5
NaYF ₄ :Yb,Tm	TiO ₂	simulated sunlight	deoxynivalenol	90.7%/60	6
NaYF ₄	SnO ₂ , Ag	simulated sunlight	RhB	100%/30	7
NaYF ₄ :Yb,Tm, Gd	NH ₂ -MIL- 101(Cr)	simulated sunlight	RhB	79%/90	This work

References

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