Electronic Supporting Information for:

Tuning the Upconversion Luminescence of Cubic KMnF₃:Yb³⁺/Er³⁺

Nanocrystals through Inert Lanthanide Ions Doping

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Experimental section

Materials

YbCl₃·6H₂O (99.99%), ErCl₃·6H₂O (99.99%), GdCl₃·6H₂O (99.99%), YCl₃·6H₂O (99.99%) were supplied by Beijing Founde Star Science and Technology Co., Ltd China. MnCl₂, KF, KOH, oleic acid (OA) at AR grade were obtained from Aladdin. All other chemicals were used without further purification.

Typical preparation of KMnF₃ nanoparticles

KMnF₃:Yb³⁺/Er³⁺ nanoparticles were synthesized via a facile hydrothermal method by using oleic acid as capping agent. In a typical procedure, 1.5g KOH was dissolved in 3mL H₂O, then 6 mL C₂H₅OH and 9 mL OA were mixed together to form a homogeneous viscous solution. A mixed aqueous solution contains 0.88 mmol MnCl₂, 0.10 mmol YbCl₃, 0.02 mmol ErCl₃ was subsequently added under intense stirring. 30 mins later, 5 mmol KF aqueous solution was ransferred to a 50 mL Teflon-lined autoclave, and then kept at 180 °C for 24h. When the reaction is over, the autoclave was cooling down to room temperature, and the products were obtained through centrifugation and washed with ethanol for several times.

Characterizations

The morphology and size of the resulting nanoparticles were investigated by means of field emission scanning electron microscope (FESEM, MX2600FE, AIKE Sepp, Oxford, UK). The powder X-Ray diffraction (XRD) pattern was carried out on a Rigaku D/max- γ B diffractometer equipped with a rotating anode and a Cu K α source (λ =0.15418 nm). The 2 θ angle of the XRD

spectra was recorded at a scanning rate of 5 °/min from 10° to 80°. The upconversion luminescence spectra were recorded using a lens-coupled monochromator (Zolix Instruments Co. Ltd, Beijing, China) of 1 nm spectral resolution with an attached photomultiplier tube (Hamamatsu CR131, Hamamatsu Photonics, Hamamatsu, Japan). An 842 nm short pass filter was placed before the spectrometer to cut off the excitation light.



Figure S1. (a) and (b) the enlarged XRD patterns comparation of $KMnF_3:10Yb^{3+}/2Er^{3+}$ nanocrystals doped with 0% and 25% Gd³⁺ ions.



Figure S2. Fourier transform infrared (FTIR) patterns of KMnF₃:10Yb³⁺/2Er³⁺ nanoparticles. The surfactant layer around the nanoparticles is identified with Fourier transform infrared (FTIR) analysis. The band at 3450 cm⁻¹ arises from the O-H (COO-H) stretching vibration. The transmission bands at 2930 and 2858 cm⁻¹ are appreciable for the asymmetric (vas) and symmetric (vs) stretching vibration of methylene (CH₂). The peaks at 3007 cm⁻¹ and 1705 cm⁻¹ are assigned to the dC-H stretching vibration and CdO stretching vibration frequency, respectively. The bands at 1564 and 1463 cm⁻¹ can correspond to the asymmetric (vas) and symmetric (vs) stretching vibration of the carboxylic group (-COOH). This result suggests the

presence of a certain amount of oleic acid on the surface of cubic nanoparticles.

Point	samples –	CIE chromaticity coordinate	
		X	У
а	KMnF ₃ :10Yb ³⁺ /2Er ³⁺	0.6621	0.2847
b	$KMnF_3:10Yb^{3+}/2Er^{3+}/5Gd^{3+}$	0.6126	0.3256
c	$KMnF_3:10Yb^{3+}/2Er^{3+}/10Gd^{3+}$	0.4943	0.4402
d	$KMnF_3:10Yb^{3+}/2Er^{3+}/15Gd^{3+}$	0.4756	0.4790
e	$KMnF_3:10Yb^{3+}/2Er^{3+}/20Gd^{3+}$	0.4203	0.4996
f	$KMnF_3:10Yb^{3+}/2Er^{3+}/25Gd^{3+}$	0.3332	0.6211



Figure S3. UC emission spectra (a) and G/R ratio (b) of $KMnF_3:10Yb^{3+}/2Er^{3+}$ with different Y^{3+} doping concentrations.



Figure S4. Decays of red upconversion emission at 660nm in $KMnF_3$:10Yb³⁺/2Er³⁺ nanocrystals with different concentration of Gd³⁺ ions: 0% to 25% under the excitation of 980 nm laser.

Table S2. Cell edges and agreement factors of $KMnF_3:10Yb^{3+}/2Er^{3+}$ with different Gd^{3+} doping concentrations by XRD define.

Sample	a/b/c (Å)	Cell Volume (Å ³)	RF-factor
0% Gd	4.18781	73.44478	4.298
5% Gd	4.18826	73.46845	2.683
10% Gd	4.18838	73.47477	3.530
15% Gd	4.18891	73.50267	3.359
20% Gd	4.18937	73.52688	3.337
25% Gd	4.19033	73.57744	5.932