

Supporting Information

Polarization modulated upconversion luminescence: single particle vs few-particle aggregate

Ping Chen,¹ Min Song,² E Wu,² Botao Wu,² Jiajia Zhou,³ Heping Zeng,² Xiaofeng Liu,^{1,*}
Jianrong Qiu^{1,4,*}

¹ School of Materials Science and Engineering, Zhejiang University, Hangzhou 310027, China

² State Key Laboratory of Precision Spectroscopy, East China Normal University, Shanghai
200062, China

³ College of Materials Science and Engineering, China Jiliang University, Hangzhou 310018,
China

⁴ State Key Laboratory of Luminescent Materials and Devices, South China University of
Technology, Guangzhou 510640, China

Contents:

1. Experimental Section

2. Supplementary data, Figure S1-S5

1. Experimental Section

$\text{Y}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ (99.99%), $\text{Er}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ (99.99%) were purchased from Ansheng Inorganic Materials Center (Ganzhou) in China. NaOH, NH_4F , NaF, HNO_3 , sodium citrate, EDTA, ethanol, 1-octadecene (90%), and oleic acid (90%) were purchased from Sigma-Aldrich. All of the chemicals were used as starting materials without further purification.

1.1 Synthesis of NaYF_4 : 5% Er nanodisks

NaYF_4 : 5% Er nanodisks were prepared with a facile and mild hydrothermal process, in which pH value in the reaction system is critical external parameters for determining the architectural features of the β - NaYF_4 nanodisks.¹ In a typical synthesis, an aqueous solution (2.5 mL, 0.2 M) of $\text{Y}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ and $\text{Er}(\text{NO}_3)_3 \cdot 6\text{H}_2\text{O}$ (lanthanide ions molar ratio, Y/Er = 95:5) was mixed with an aqueous solution of sodium citrate (22.5 mL, 2.08 M) under stirring for 30 min to form a white solution. Subsequently, aqueous solution (10 mL) of NaF (10 mL, 0.625 M) was added and stirred for 1 h, resulting in a complex with the lanthanide/ sodium citrate/ NaF molar ratio being 1/90/12. Then, HNO_3 was added to adjust the pH of the solution to 3. The obtained solution was then transferred into a 50 mL autoclave and hydrothermally treated at 180 °C for 2 h. After cooling to room temperature, products were separated by centrifugation and washed with ethanol for three times and then dried at 60 °C in vacuum.

1.2 Structural characterizations

X-ray diffraction (XRD) pattern of the dry powder was obtained on a RIGAKU D/MAX 2550/PC diffractometer (Japan) with a slit of 0.02° at a scanning rate of 5° min^{-1} using $\text{Cu K}\alpha$ radiation ($\lambda = 1.5406 \text{ \AA}$). Scanning Electron Microscopy (SEM) analysis was performed on a Field Emission Scanning Electron Microscopy (FESEM) (ZEISS SU-8010). High-resolution

Transmission Electron Microscopy (HRTEM) analysis was performed on a FEG-TEM (Tecnai G2 F30 S-Twin, Philips-FEI, Netherlands) operated at 300 kV.

2. Supplementary date

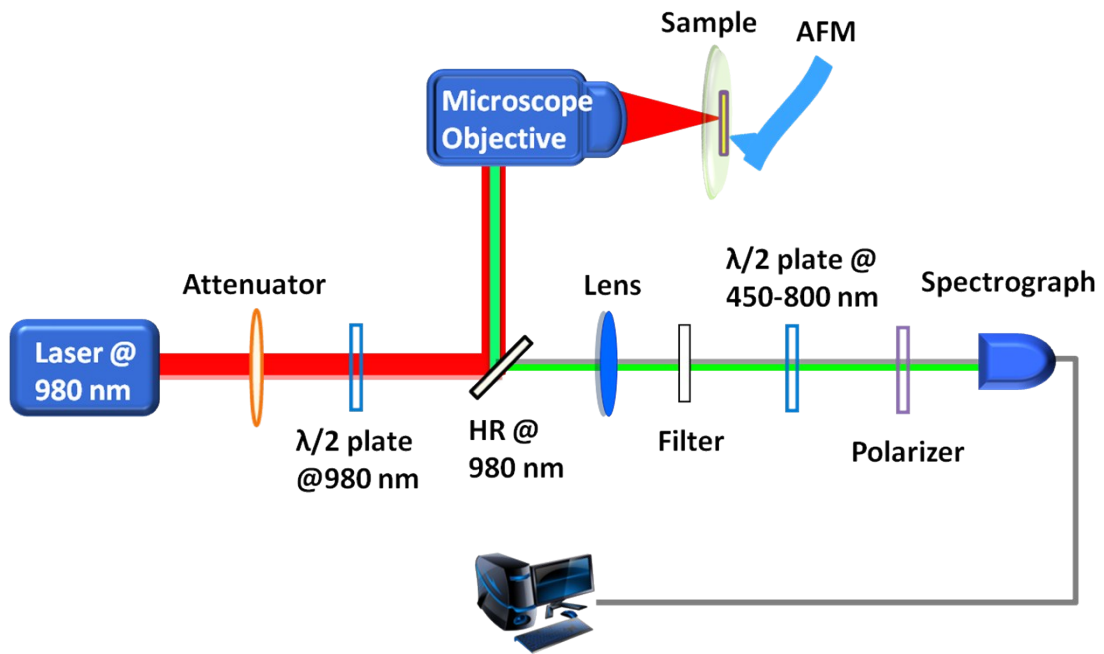


Fig. S1 Schematic diagram of luminescence spectrum test system.

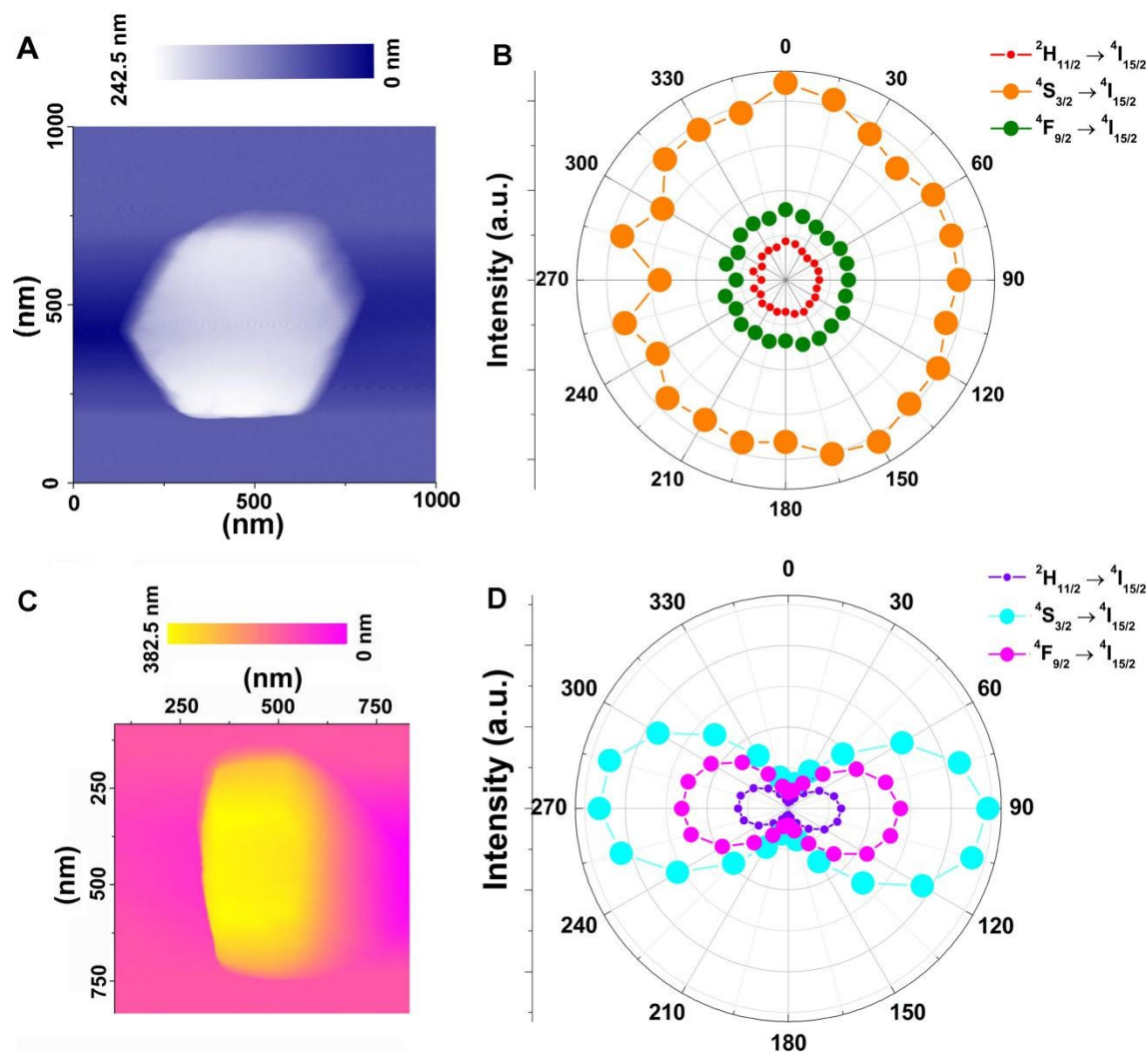


Fig. S2 A) AFM images of single nanodisk recorded for a single nanodisk in which the a axis is parallel to the horizontal plane (HP). B) Polar plots of integrated UC luminescence intensity of single nanodisk as a function of excitation polarization angle for the transitions from ${}^2\text{H}_{11/2} \rightarrow {}^4\text{I}_{15/2}$, ${}^4\text{S}_{3/2} \rightarrow {}^4\text{I}_{15/2}$, ${}^4\text{F}_{9/2} \rightarrow {}^4\text{I}_{15/2}$ of Er^{3+} , recorded at $a \parallel \text{HP}$ and $E_{ex} \parallel a$ axis @ $\theta = 0^\circ$. C) AFM images of single nanodisk recorded for a single nanodisk in which the c axis is parallel to HP. D) Polar plots of integrated UC luminescence intensity of single nanodisk as a function of excitation polarization angle for the transitions from ${}^2\text{H}_{11/2} \rightarrow {}^4\text{I}_{15/2}$, ${}^4\text{S}_{3/2} \rightarrow {}^4\text{I}_{15/2}$, ${}^4\text{F}_{9/2} \rightarrow {}^4\text{I}_{15/2}$ of Er^{3+} , recorded at $c \parallel \text{HP}$ and $E_{ex} \parallel c$ axis @ $\theta = 0^\circ$.

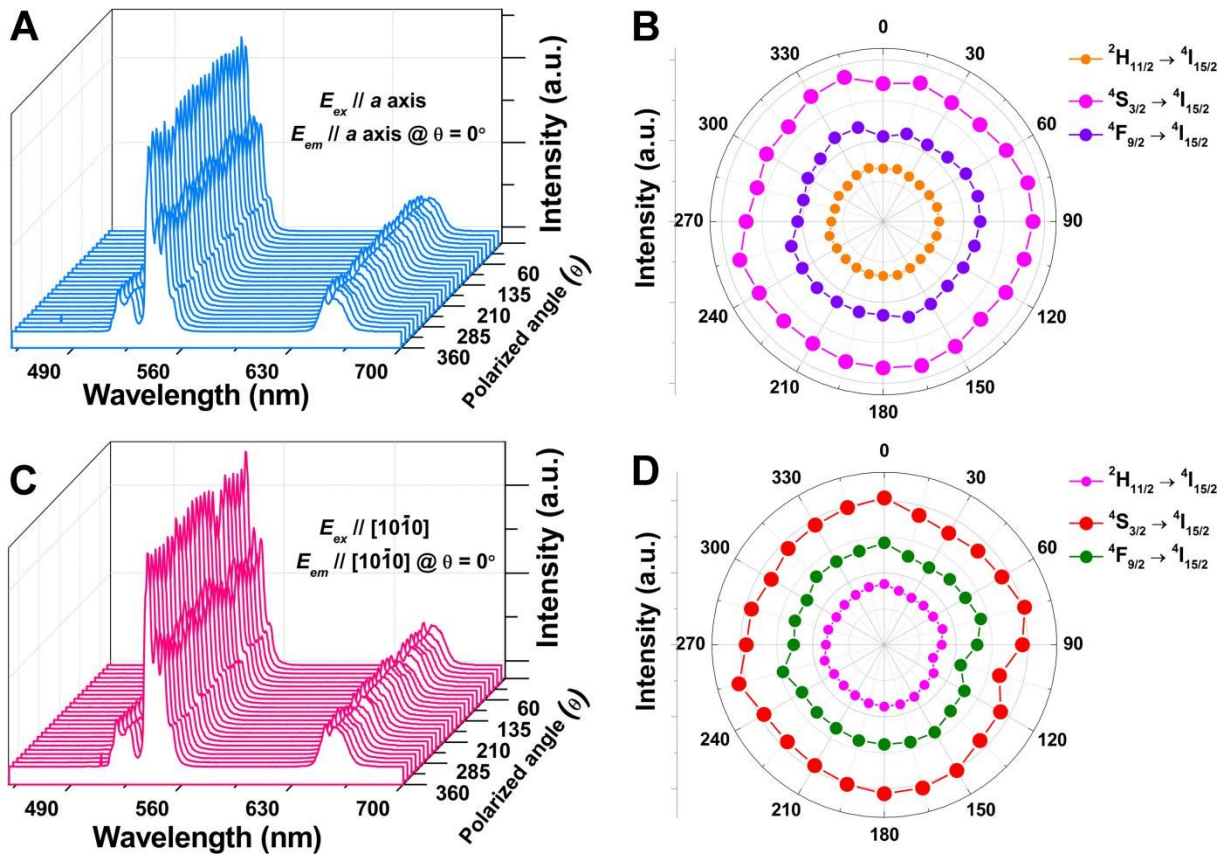


Fig. S3 A) UC luminescence spectra recorded at emission polarization angles of $0^\circ - 360^\circ$ for single nanodisk under conditions of $a \parallel \text{HP}$ and $E_{ex} \parallel a$. B) Polar plots of integrated UC luminescence intensity as a function of emission polarization angle for the transitions from ${}^2\text{H}_{11/2} \rightarrow {}^4\text{I}_{15/2}$, ${}^4\text{S}_{3/2} \rightarrow {}^4\text{I}_{15/2}$, ${}^4\text{F}_{9/2} \rightarrow {}^4\text{I}_{15/2}$ of Er^{3+} for single nanodisk under configurations of $a \parallel \text{HP}$ and $E_{ex} \parallel a$. C) UC luminescence spectra recorded at emission polarization angles of $0^\circ - 360^\circ$ for single nanodisk under conditions of $a \parallel \text{HP}$ and $E_{ex} \parallel [10\bar{1}0]$. D) Polar plots of integrated UC luminescence intensity as a function of emission polarization angle for the transitions from ${}^2\text{H}_{11/2} \rightarrow {}^4\text{I}_{15/2}$, ${}^4\text{S}_{3/2} \rightarrow {}^4\text{I}_{15/2}$, ${}^4\text{F}_{9/2} \rightarrow {}^4\text{I}_{15/2}$ of Er^{3+} for single nanodisk under configurations of $a \parallel \text{HP}$ and $E_{ex} \parallel [10\bar{1}0]$.

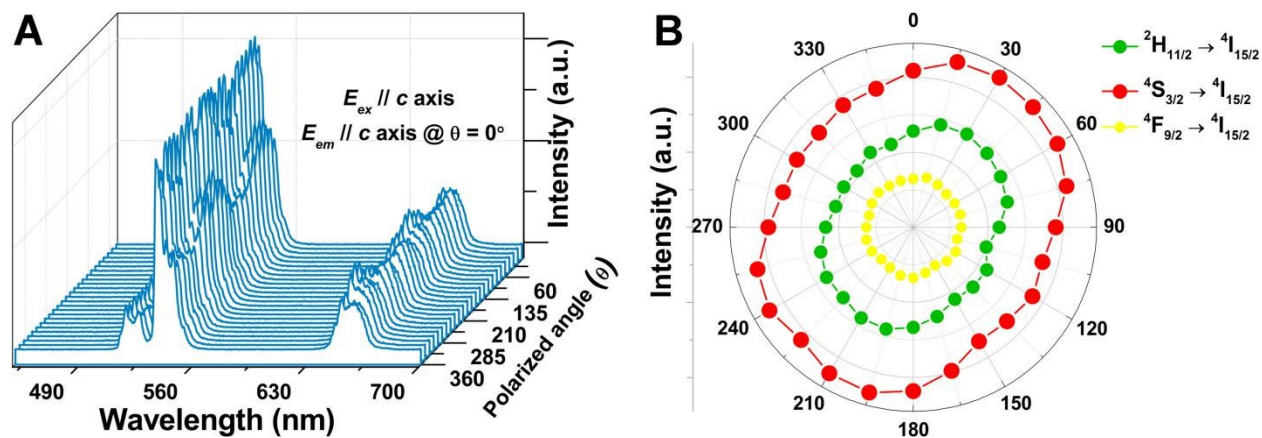


Fig. S4 A) UC luminescence spectra recorded at emission polarization angles of $0^\circ - 360^\circ$ for single nanodisk under conditions of $c // HP$ and $E_{ex} // c$. B) Polar plots of integrated UC luminescence intensity as a function of emission polarization angle for the transitions from ${}^2H_{11/2} \rightarrow {}^4I_{15/2}$, ${}^4S_{3/2} \rightarrow {}^4I_{15/2}$, ${}^4F_{9/2} \rightarrow {}^4I_{15/2}$ of Er^{3+} for single nanodisk under configurations of $c // HP$ and $E_{ex} // c$.

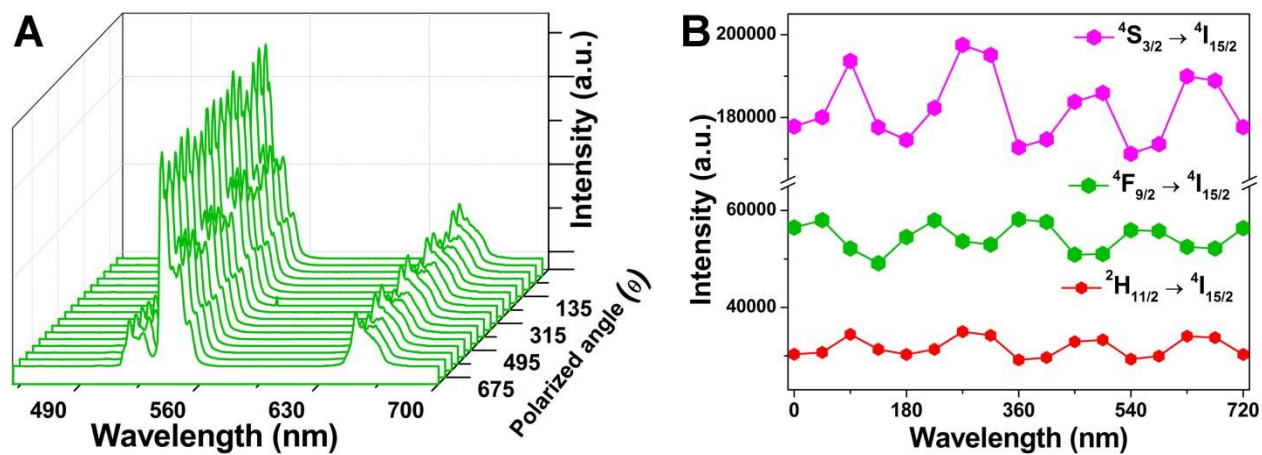


Fig. S5 A) UC luminescence spectra of the aggregated nanodisks recorded under emission polarization angle varied from 0° to 360° . C) The dependence of UC luminescence intensity of the aggregated nanodisks with different transitions of Er^{3+} on the emission polarization angle.

Reference

1. L. Liang, Y. Liu, C. Bu, K. Guo, W. Sun, N. Huang, T. Peng, B. Sebo, M. Pan, W. Liu, *et. al.* *Adv. Mater.*, 2013, **25**, 2174.