



Cite this: *Nanoscale*, 2022, **14**, 4595

Simultaneous ultraviolet-C and near-infrared enhancement in heterogeneous lanthanide nanocrystals†

Yachong Liu,^{‡,a} Mingzhu Zhou,^{‡,a} Meng-Tao Zhou,^a Han-Lin Wei,^a Yan Su^b and Qianqian Su^{†,a*}

Lanthanide-doped nanocrystals that simultaneously convert near-infrared (NIR) irradiation into emission of shorter (ultraviolet-C, UVC) and longer wavelengths (NIR) offer many exciting opportunities for application in drug release, photodynamic therapy, deep-tissue bioimaging, and solid-state lasing. However, a formidable challenge is the development of lanthanide-doped nanocrystals with efficient UVC and NIR emissions simultaneously due to their low conversion efficiency. Here, we report a dye-sensitized heterogeneous core–multishell architecture with enhanced UVC emission and NIR emission under 793 nm excitation. This nanocrystal design efficiently suppresses energy trapping induced by interior lattice defects and promotes upconverted UVC emission from Gd^{3+} . Moreover, a significant downshifting emission from Yb^{3+} at 980 nm was also observed owing to an efficient energy transfer from Nd^{3+} to Yb^{3+} . Furthermore, by taking advantage of ICG sensitization, we realized a largely enhanced emission from the UVC to NIR spectral region. This study provides a mechanistic understanding of the upconversion and downshifting processes within a heterogeneous architecture while offering exciting opportunities for important biological and energy applications.

Received 5th November 2021,
Accepted 23rd February 2022

DOI: 10.1039/d1nr07329j

rsc.li/nanoscale

Introduction

The applications of lanthanide luminescence in bioimaging, theranostics, displays, photovoltaics, and in the emerging field of optogenetics have driven the growing demand for rational control over the emission profiles of lanthanide-doped nanocrystals.^{1–12} Over the past decade, the toolbox developed for the generation of efficient upconversion emission includes mechanistic strategies such as excited state absorption (ESA), energy transfer upconversion (ETU), energy migration upconversion (EMU), and interfacial energy transfer (IET).^{13–16} These strategies combined with nanostructure engineering enabled a wide range of upconversion activators with emissions ranging from ultraviolet (UV) to near-infrared (NIR).^{17,18}

Despite the enriched emission profiles, their practical implementation in various applications is hindered by the limited emission intensity and the difficulty in shifting the

emission wavelength to the ultraviolet-C (UVC, 100–290 nm) region. This is because photon upconversion from high-lying energy levels is significantly influenced by many deleterious factors, such as cross-relaxation between lanthanide ions, concentration quenching, and energy consumption induced by interior lattice defects.^{19–21} Recently, we proposed an upconverted excitation lock-in (UCEL) strategy to enhance UVC emission in a Nd^{3+} -sensitized heterogeneous nanostructure. Our study showed that efficient UVC emission and six-photon upconversion emission at 253 nm under 808 nm excitation are possible *via* embedding a $NaYF_4$ interlayer. The $NaGdF_4@NaYF_4@NaGdF_4@NaGdF_4$ heterogeneous core–multishell structure preserves the excitation energy within the core domain and effectively restrains energy dissipation by interior lattice defects.²² However, our previous nanostructural design still suffers from the limited conversion of the excitation irradiation to radiative UVC emission owing to the small absorption cross-section of lanthanides.²³

Concurrently, lanthanide-based NIR (700–1700 nm) imaging probes are ideal candidates for biomedical imaging because of minimal light absorption, reduced light scattering, and negligible autofluorescence. These probes could monitor the physiological processes of deep tissues with high imaging resolution.^{24–33} Therefore, a variety of NIR nanophosphors have been developed, such as rare earth fluorides, rare earth

^aInstitute of Nanochemistry and Nanobiology, Shanghai University, Shanghai 200444, China. E-mail: chmsqq@shu.edu.cn

^bGenome Institute of Singapore, Agency of Science Technology and Research, 138672, Singapore

†Electronic supplementary information (ESI) available. See DOI: 10.1039/d1nr07329j

‡These authors made an equal contribution to this work.

vanadate, rare earth oxides nanophosphors, *etc.* However, the vast majority of researchers have only focused on the optical studies of downshifting nanocrystals with emission in the NIR region.^{34–36} Noteworthy, the simultaneous generation of short-wavelength (UVC) and long-wavelength (NIR) light by one nanocrystal may offer many exciting opportunities for application in deep-tissue NIR imaging guided drug release, photodynamic therapy, and more. However, little effort has been made to develop nanocrystals that simultaneously give rise to UVC and NIR luminescence.

Herein, we introduce a rational heterogeneous core–multi-shell strategy that provides precise control over the upconverted excitation energy flux (Scheme 1). In our design, a large amount of Nd³⁺ ions (90%) is selectively doped within the second shell layer for effective harvesting of light with wavelengths around 793 nm. The designed nanostructure with a Gd-based core and a Y-based shell preserves the upconverted UVC energy within the core domain. The effective energy transfer from Nd³⁺ to Yb³⁺ ions enables NIR downshifting emission at around 980 nm. The organic dye ICG acts as an antenna to absorb incident light and transfer the excited energy to our designed nanocrystals, significantly improving the emission intensity of lanthanides from the UVC to NIR region. This study implies that Nd³⁺-sensitized heterogeneous core–multi-shell nanocrystals with a new emission profile offer more opportunities for designing efficient upconversion nanoplat-forms in various fundamental research and technological applications.

Results and discussion

Design of Nd³⁺-doped nanocrystals

In our study, we designed a heterogeneous core–multishell nanocrystal to suppress interior energy consumption and surface quenching, achieving tunable UVC and NIR emissions. Hexagonal phase NaGdF₄:49%Yb,1%Tm was chosen as a core nanocrystal for its ability to render the possibility of UVC emission.²² The first shell layer of NaYF₄:20%Yb is optically inert to

the excited states of Gd³⁺ ions and can block the energy transfer from Gd³⁺ to interior lattice defects. The second shell layer contained a high concentration of Nd³⁺ ions (90 mol%) for harvesting light. A 793 nm diode laser was selected as the excitation source because the absorption intensity at 793 nm for the nanocrystals is significantly higher than that at 808 nm and 980 nm. The approximate absorption cross-section σ of Nd³⁺ at 793 nm, 808 nm and 980 nm was calculated to be $4.3 \times 10^{-19} \text{ cm}^2$, $1.3 \times 10^{-19} \text{ cm}^2$ and $4.3 \times 10^{-20} \text{ cm}^2$ from the UV-vis absorption spectra of the nanocrystals (Fig. S1†).³⁷

Synthesis and characterization of Nd³⁺-doped GYNY nanocrystals

In a typical experiment, we synthesized NaGdF₄:49%Yb,1%Tm@NaYF₄:20%Yb@NaNdF₄:10%Yb@NaYF₄ (GYNY) heterogeneous core–multishell nanocrystals *via* a modified literature method (Fig. 1a).^{22,38} As shown in Fig. S2,† the size of NaGdF₄:49%Yb,1%Tm was found to be 8.6 nm. The transmission electron microscopy (TEM) images show that each shell layer is about 1.5–2.5 nm in thickness and the obtained GYNY nanocrystals are about 21 nm on average. High-resolution TEM (HR-TEM) indicates a single crystalline structure of the as-synthesized GYNY nanocrystals, and the lattice distance of 0.52 nm corresponds to the *d* spacing for the (100) plane of hexagonal NaYF₄ (Fig. 1b). Energy-dispersive X-ray spectroscopy confirms the presence of Nd³⁺, Gd³⁺, Yb³⁺, Y³⁺ and Tm³⁺ in these GYNY nanocrystals (Fig. S3†). X-ray powder diffraction (XRD) confirmed the hexagonal phase of the as-prepared nanocrystals (JCPDS file number 16-0334, Fig. S4†). We have also conducted elemental mapping and energy dispersive X-ray (EDX) line scan on the whole particle to demonstrate the spatial distribution of rare-earth cations and the heterogeneous core–shell structure. As shown in Fig. 1b, Nd and Y ions are concentrated in the periphery region, while Gd and Yb show apparently higher concentrations in the central region. The relative concentrations of Nd, Y, Gd and Yb along the scanning path are plotted as shown in Fig. S5.† The result provides more signal counts of Gd and Yb from the central region while those of Nd and Y from the periphery region. Of note is that our nanoparticles do not show a sharp interface at the atomic scale. This result could be ascribed to core–shell intermixing caused by the relatively small core (8.6 nm) and whole core–multishell nanoparticle (20.6 nm), which is consistent with the previous report.^{39–44}

We conducted a group of experiments to compare the emission intensity of GYNY nanocrystals under 793 nm, 808 nm and 980 nm excitation. As expected, GYNY nanocrystals under 793 nm excitation showed the strongest emission compared to those under 808 nm and 980 nm excitation (Fig. 1c), which is consistent with our absorption cross-section results of Nd³⁺ at 793 nm, 808 nm and 980 nm. Similarly, the comparison of the downshifting emission intensities at 793 nm, 808 nm and 980 nm showed the same trend (Fig. S6†).

We next studied the photoluminescence spectrum of the as-synthesized GYNY nanocrystals. As shown in Fig. 1d, obvious upconversion emissions from the ⁶I₇ and ⁶P₇ energy levels of



Scheme 1 Schematic illustration of the dye-sensitized upconversion and downshifting mechanism for UVC and NIR generation.

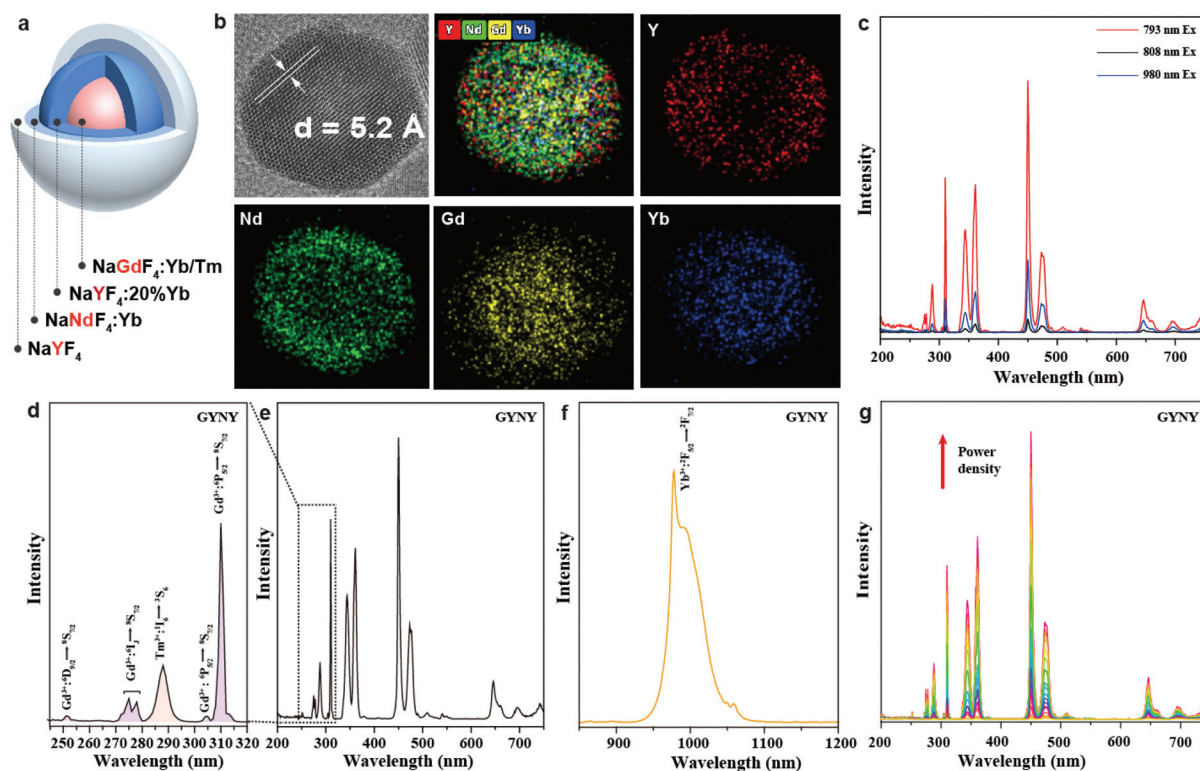


Fig. 1 (a) Schematic illustration of a NaGdF₄:49%Yb,1%Tm@NaYF₄:20%Yb@NaNdF₄:10%Yb@NaYF₄ (GYNY) nanoparticle. (b) High-resolution TEM image and elemental mapping of the corresponding GYNY nanocrystal. (c) Corresponding upconversion luminescence spectra of GYNY nanoparticles in cyclohexane under 793, 808 and 980 nm excitation at a power density of 28.0 W cm⁻². (d–f) Room-temperature luminescence spectra in the range of 240–750 nm and 850–1200 nm of GYNY nanoparticles in cyclohexane. (g) Excitation-power-dependent upconversion emission spectra of GYNY nanoparticles under 793 nm excitation.

Gd³⁺ peaked at 273 nm (⁶I_{7/2} → ⁸S_{7/2}), 276 nm (⁶I_{7/2} → ⁸S_{7/2}), 279 nm (⁶I_{7/2} → ⁸S_{7/2}), 306 nm (⁶P_{5/2} → ⁸S_{7/2}) and 311 nm (⁶P_{7/2} → ⁸S_{7/2}) in the UV region were observed under 793 nm excitation. A small peak at 253 nm from the ⁶D_J energy levels of Gd³⁺ demonstrated a six-photon upconversion process, which could be attributed to the ⁶D_{9/2} → ⁸S_{7/2} optical transition.^{45–47}

Meanwhile, we observed an intense downshifting emission peaked at 980 nm (²F_{5/2} → ²F_{7/2}) from Yb³⁺ due to the efficient energy transfer from Nd³⁺ to Yb³⁺ in GYNY nanocrystals (Fig. 1f). We further investigated the excitation-power-dependent upconversion emission spectra of GYNY nanoparticles under 793 nm excitation (Fig. 1g). Taken together, photoluminescence investigation revealed that a collection of lanthanide ions including Gd³⁺, Tm³⁺, Nd³⁺ and Yb³⁺ can work together to generate upconversion emissions spanning the UVC and NIR spectral region from 250 to 1200 nm.

The generation of a broad emission profile spanning the whole UV to NIR spectral region is important for expanding the scope of lanthanide-based nanocrystals in favor of diverse applications in bioimaging, molecular sensing, therapy, and security printing. Therefore, on the basis of this nanocrystal design strategy, we further broaden the spectrum of lanthanide-based nanocrystals by simply doping Yb and Er in the third shell layer. We prepared NaGdF₄:49%Yb,1%

Tm@NaYF₄:20%Yb@NaNdF₄:10%Yb@NaYF₄:20%Yb,2%Er@NaYF₄ (GYNY_{20%Yb,2%Er}Y) nanocrystals by the same synthesis route except for the use of the third shell stock solution of NaYF₄:Yb,Er. Significantly, the luminescence spectrum of the samples could extend to 1600 nm (Fig. 2).

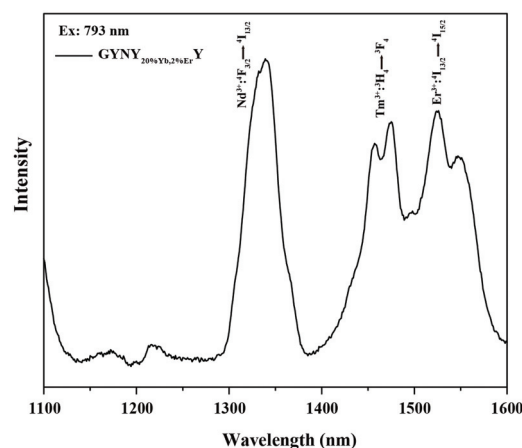


Fig. 2 Downshifting emission spectrum of NaGdF₄:49%Yb,1%Tm@NaYF₄:20%Yb@NaNdF₄:10%Yb@NaYF₄:20%Yb,2%Er@NaYF₄ nanocrystals.

Optimization of dopant concentration

To achieve the required strong emission in both the UVC and NIR regions in a nanocrystal, we synthesized a series of NaGdF₄@NaYF₄@NaNdF₄@NaYF₄ core-multishell nanocrystals with varying contents of Yb³⁺ and Nd³⁺. The second shell layer was doped with different concentrations of Yb³⁺ (0–50 mol%), while it had a fixed Nd³⁺ dopant concentration of 50 mol% (Fig. S7a and S8†). We then fixed the Yb³⁺ dopant concentration of 10 mol% and changed the concentration of Nd³⁺ from 1 mol% to 90 mol% in the second shell layer (Fig. S7b and S9†). Taking into account the UVC upconversion and downshifting emission intensity, we determined the optimal Yb³⁺ and Nd³⁺ concentration to be 10 and 90 mol%, respectively (Fig. S7b and S7c†).

The role of the NaYF₄ interlayer in GYNY nanocrystals

To verify the role of the NaYF₄-based first layer in realizing UVC upconversion emission, we synthesized a pair of GYNY and NaGdF₄:49%Yb,1%Tm@NaGdF₄:20%Yb@NaNdF₄:10%Yb@NaYF₄ (GGNY) nanocrystals. After replacing the interlayer of NaYF₄ with NaGdF₄, the upconversion emissions from ⁶D_J, ⁶I_J, and ⁶P_J in the UV region were significantly quenched either under 793 nm or 980 nm excitation (Fig. 3a and S10†). We then compared the excited state lifetime of

Gd³⁺. As shown in Fig. 3b–e, a significantly prolonged Gd³⁺ lifetime (⁶P_{7/2} level, ~6 times) was observed when the NaYF₄ first layer was applied. In stark contrast, the Gd³⁺ lifetimes for emissions from the ⁶D_J and ⁶I_J energy levels and the Tm³⁺ lifetimes for emissions from the ¹I₆ energy level were essentially unaltered after the use of the NaYF₄ first layer. These results indicated that the energy loss from Gd³⁺ to interior lattice defects was mainly through the ⁶P_{7/2} energy level.

Of note is that real materials usually contain a high density of lattice defects. An excited Gd³⁺ ion undergoes two different optical processes: (i) an excited Gd³⁺ ion transfers its energy to a nearby defect centre and subsequently undergoes non-radiative relaxation; (ii) a resonant energy transfer occurs within the Gd³⁺ sublattice, followed by radiative relaxation and gives rise to upconversion emission. As shown in Fig. 3f, the NaYF₄-based first shell can block the energy transfer from Gd³⁺ to the interior lattice defects located in the shell regions in our design, promoting the occurrence of fast energy hopping in the Gd³⁺ sublattice at the core domain, thereby realizing intense UVC upconversion emissions. The NaYF₄-based first layer played a critical role in locking in and recycling Gd³⁺ excitation energy. These results accord with the recently proposed upconverted excitation lock-in (UCEL) mechanism by our group.²²



Fig. 3 (a) Upconversion emission spectra of GYNY and GGNY nanocrystals under 793 nm excitation. (b and c) Upconversion luminescence decay curves of Gd³⁺ emissions at 253 nm and 276 nm in GYNY and GGNY by pulsed 793 nm excitation. (d and e) Upconversion luminescence decay curves of Tm³⁺ emissions at 290 nm and Gd³⁺ emission at 311 nm in GYNY and GGNY by pulsed 793 nm excitation. (f) Schematic illustration of the energy transfer mechanism for UVC and NIR generation within the GYNY heterogeneous nanocrystal.

The energy transfer efficiency from Nd³⁺ to Yb³⁺

The energy transfer efficiency η from Nd³⁺ to Yb³⁺ can be quantitatively estimated from eqn (1) and (2):^{48,49}

$$\eta = 1 - \frac{\tau_m}{\tau_{Nd}} \quad (1)$$

$$\tau_m = \frac{\sum \alpha_i \tau_i^2}{\sum \alpha_i \tau_i} \quad (2)$$

where τ_m is the mean lifetime of the energy donor Nd³⁺ in the presence of the energy acceptor Yb³⁺, τ_{Nd} is the intrinsic lifetime of Nd³⁺, and α is the amplitude.

To calculate the Nd³⁺-to-Yb³⁺ energy transfer efficiency, we synthesized a pair of nanocrystals of NaGdF₄:49%Yb,1%Tm@NaYF₄:20%Yb@NaNdF₄:10%Yb@NaYF₄ (GYNY with Yb³⁺) and NaGdF₄:49%Yb,1%Tm@NaYF₄@NaNdF₄:10%Yb@NaYF₄ (GYNY without Yb³⁺) as shown in Fig. S11.† Of note is that the energy acceptor of Yb³⁺ was replaced by optically inert Y³⁺ ions to measure the intrinsic lifetime of Nd³⁺. Under 793 nm pulsed excitation, the lifetime of Nd³⁺ of the pair of nanocrystals at 893 nm was measured, and the Nd³⁺-to-Yb³⁺ energy transfer efficiency was calculated to be 89% (Fig. 4), which is slightly higher than the previous reports.⁵⁰

Dye sensitization for upconversion and downshifting luminescence enhancement

For efficient dye sensitization, the absorption wavelength of the lanthanide-based nanocrystals should overlap with the dye fluorescence to achieve an efficient energy transfer from the dye to nanocrystals. Herein, indocyanine green (ICG) was chosen for NIR dye sensitization due to its large absorption cross-section ($\sim 6 \times 10^{-16}$ cm²), which is about 1500 times higher than that of Nd³⁺ at 793 nm ($\sim 4 \times 10^{-19}$). The absorption curve of ICG was recorded using a UV-vis spectrophotometer. As shown in Fig. 5a, a good spectral overlap between

the absorption band of GYNY and the fluorescence of ICG was observed, providing an important prerequisite for efficient non-radiative energy transfer from ICG to GYNY nanocrystals.⁵¹

We next investigated the optical behavior of GYNY nanocrystals sensitized by the ICG dye of different concentrations. To ensure close contact between nanocrystals and ICG, the original oleic acid ligand capped on the surface of GYNY was replaced by the ionic NOBF₄ ligand, and then the ICG dye was coated onto the surface of nanocrystals.⁵² The removal of oleic acid and the surface capping of ICG were confirmed by FTIR analysis (Fig. S12†). After capping ICG on the surface of GYNY, we observed a remarkably enhanced absorption peak of GYNY, which also confirmed the success of ICG coating (Fig. 5b).

Subsequently, we found that the enhancement factor varies with changes in ICG concentration (Fig. S13†). In the tested ICG concentration range, the intensities of upconversion and downshifting emissions of GYNY were all significantly improved, reaching a maximum at the ICG concentration of 100 μ L. Specifically, we observed more than 30-fold, 10-fold, and 8-fold enhancements in the range of 200–400 nm of UVC emission, 400–700 nm of visible emission, and 850–1200 nm of NIR emission by dye sensitization (Fig. 5c–e). The enhancement factor of the whole spectrum ranging from 240 to 1200 nm of the as-prepared ICG sensitized-GYNY compared with that of GYNY was estimated to be 8.6-fold. Notably, a higher concentration of ICG induced a small emission enhancement factor, which is possibly due to the dye–dye quenching interaction on the surface of GYNY.⁵² The photograph of ICG sensitized-GYNY showed an obviously increased brightness, providing direct evidence of effective ICG sensitization for Nd³⁺-doped nanocrystals under 793 nm excitation (Fig. 5e, insert).

We also extended the strategy by coating the ICG dye on a novel nanoparticle structure, NaGdF₄:49%Yb,1%Tm@NaYF₄:20%Yb@NaNdF₄:10%Yb@NaYF₄:20%Yb,2%Er@NaYF₄ (GYNY_{20%Yb,2%Er}). The results showed that the luminescence intensity of Tm upconversion and Er downshifting can be enhanced simultaneously under 793 nm excitation. We observed about 2.4-fold and 2.1-fold enhancements in the range of 200–700 nm and 1250–1600 nm by dye sensitization, respectively (Fig. S14†).

Notably, ICG is a tricarboxyanine and fluorescent dye, which will undergo photodegradation and thermal degradation under certain conditions.^{53,54} We studied the photostability of GYNY@ICG under excitation at 793 nm. The emission spectra of GYNY@ICG were obtained every 2 min at a power density of 20 W cm⁻². Fig. S15† shows that the Förster resonance energy transfer (FRET) effect between the dye and Nd decreased with a prolonged illumination time and disappeared at about 20 min.

Moreover, we conducted a quantitative study for the quantum yields (QYs) of GYNY and GYNY@ICG nanoparticles. We measured the QYs of upconversion emission in the range of 200–400 nm of the as-prepared GYNY and GYNY@ICG nanoparticles, and the results were recorded to be approxi-

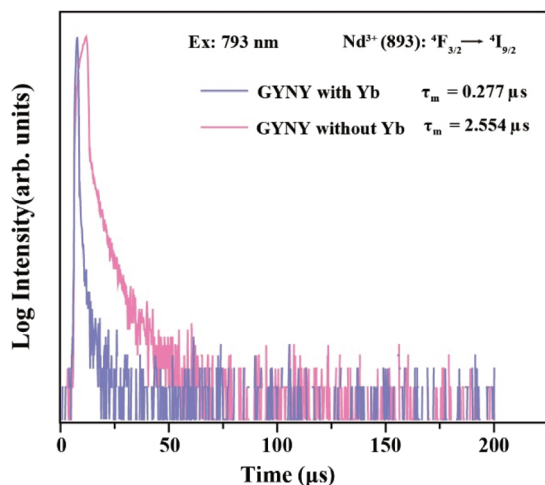


Fig. 4 Luminescence decay curves of Nd³⁺ emissions measured at 893 nm for GYNY with and without Yb³⁺ nanocrystals by pulsed 793 nm excitation.

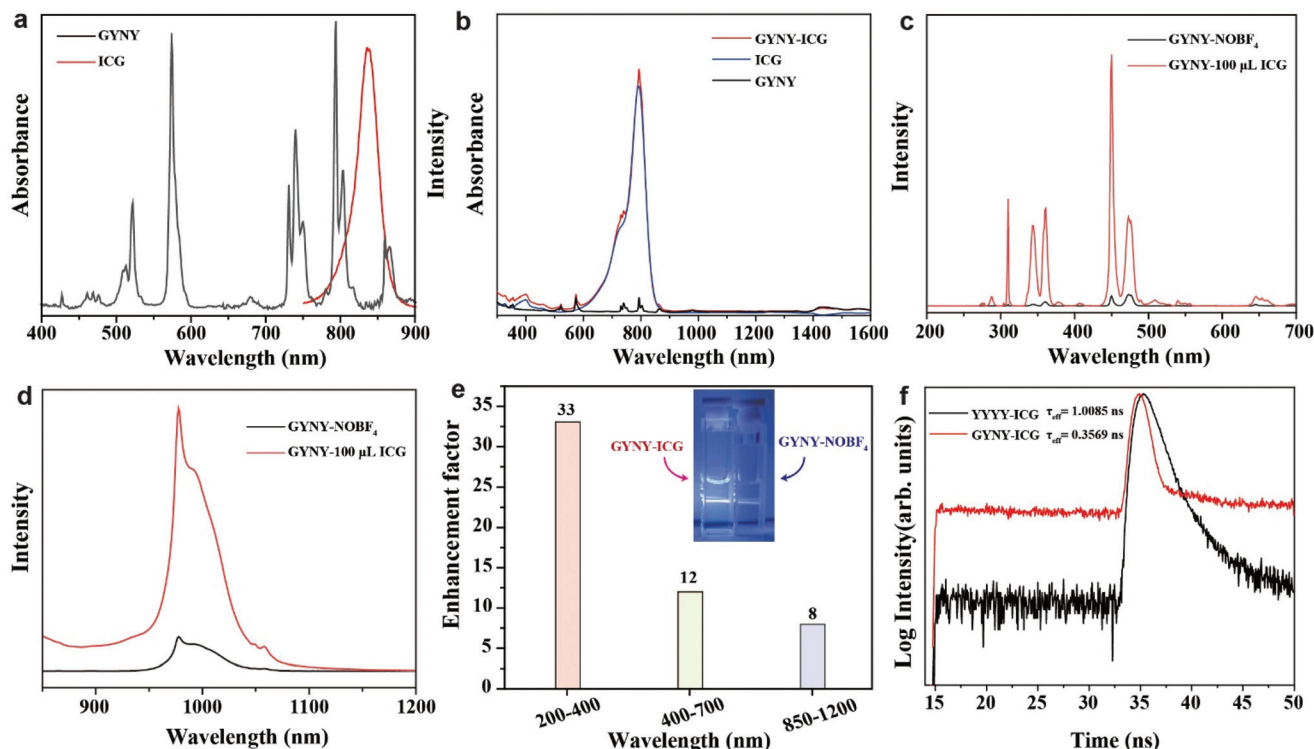


Fig. 5 (a) Absorption spectrum of GYNY nanocrystals (black line) and the fluorescence spectrum of ICG (red line) under 730 nm excitation. (b) Absorption spectra of GYNY, ICG and ICG-coated GYNY nanocrystals; the concentrations of GYNY and ICG were 5 mg mL⁻¹ and 5 μg mL⁻¹, respectively. (c and d) Upconversion and downshifting emission of GYNY and ICG-coated GYNY nanocrystals at the optimized concentration. (e) Enhancement factor of the luminescence of GYNY nanocrystals at 200–400 nm, 400–700 nm, 850–1200 nm after ICG coating. (f) Luminescence decay curves of ICG fluorescence obtained at 830 nm for GYNY and YYYI nanocrystals with ICG coating by pulsed 270 nm excitation.

mately 0.04% and 0.14%, respectively. Besides, the QYs of GYNY and GYNY@ICG nanoparticles were approximately 0.45% and 0.63% from 400 to 750 nm, 0.49% and 0.87% from 200 to 750 nm, and 14.88% and 14.97% from 820 to 1400 nm, respectively.

We then obtained the fluorescence decay curves of the ICG emission at 830 nm under 270 nm excitation. As shown in Fig. 5f, the decay lifetimes of the ICG dye with and without the presence of GYNY nanocrystals were determined to be 0.3569 ns and 1.0085 ns, respectively, suggesting an energy transfer efficiency of 64.6%. This result also demonstrated a Förster resonance energy transfer mechanism from ICG to Nd³⁺ ions doped in the nanocrystals.⁵⁵

In our study, the critical distance from the dye to the Nd ions was determined by adjusting the thickness of the outermost NaYF₄ layer to 2.3 nm (GYNY), 4.8 nm (GYNYY) and 6.7 nm (GYNYYY), respectively (Fig. S16†). Then, the materials with a concentration of 5 mg mL⁻¹ were coated with the ICG dye (100 μL, 0.1 mg mL⁻¹), respectively. We compared the upconversion luminescence intensity of the nanoparticles with different thicknesses of the NaYF₄ layer before and after coating with the dye (Fig. S16†). When the thickness of the outermost NaYF₄ layer was 4.8 nm, the emission intensity was almost unchanged after coating with the ICG dye in the range of 200–700 nm, respectively. However, when the thickness of

the outermost NaYF₄ layer was 6.7 nm, the emission intensity decreased after coating with the ICG dye. These results showed that the critical distance between the ICG dye and Nd is about 5 nm.

In addition, we also investigated whether the triplet state of ICG is involved in the energy transfer between Gd. Herein, we designed a new nanostructure of GYN@Y30Gd by doping 30% Gd in the outermost NaYF₄ shell. After coating ICG onto the surface of GYN@Y30Gd and GYNY, we compared their luminescence intensities before and after ICG dye coating. The results indicated that the emission intensity of GYNY and GYN@Y30Gd from 200 to 400 nm was enhanced by about 4 and 2 times, and that from 850 to 1200 nm was enhanced by about 2-fold and 3-fold, respectively (Fig. S17a–d†). This enhancement effect was negligible compared with the huge enhancement in the triplet state of ICG involved energy transfer in the previous report.⁵⁶ Moreover, there is no significant difference in the lifetimes of Gd³⁺ (311 nm) in GYNY and GYN@Y30Gd nanoparticles before and after ICG coating (Fig. S17e†). Therefore, we can infer that there is no energy transfer between the triplet state of the ICG dye and Gd.

Considering its potential in *in vivo* and *in vitro* biological applications, it is essential to achieve high-efficiency dye-sensitized upconversion and downshifting luminescence in aqueous solutions.⁵² We connect ICG to the surface of GYNY

nides and enhancing the NIR dye-sensitized UVC to NIR emission. This study will also benefit the future development of novel lanthanide nanocrystals for applications in biomedicine, photocatalysis, environmental science, and many other emerging fields.

Experimental section

Materials, details on nanocrystal synthesis and characterization are included in the ESI.†

Author contributions

Q. S. planned the project. Y. L., M. Z., M.-T. Z., and H.-L. W. performed the experiments. Q. S., Y. L., M. Z., and Y. S. designed the experiments and analysed the data. Q. S., Y. L. and M. Z. wrote the paper and prepared the figures. All authors discussed the results and revised the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

The authors thank the National Natural Science Foundation of China (no. 21701109).

Notes and references

- Q. Su, W. Feng, D. Yang and F. Li, *Acc. Chem. Res.*, 2017, **50**, 32–40.
- M. Ang, J. Yoon, M. Zhou, H. Wei, Y. Goh, Z. Li, J. Feng, H. Wang, Q. Su, D. Ong and X. Liu, *Adv. Mater.*, 2022, **34**, 2106194.
- X. Zhu, Q. Su, W. Feng and F. Li, *Chem. Soc. Rev.*, 2017, **46**, 1025–1039.
- D. Yang, P. Ma, Z. Hou, Z. Cheng, C. Li and J. Lin, *Chem. Soc. Rev.*, 2015, **44**, 1416–1448.
- F. Wang and X. Liu, *Chem. Soc. Rev.*, 2009, **38**, 976–989.
- S. Wang, L. Liu, Y. Fan, A. M. El-Toni, M. S. Alhoshan, D. Li and F. Zhang, *Nano Lett.*, 2019, **19**, 2418–2427.
- L. Wang, H. Zhou, J. Hu, B. Huang, M. Sun, B. Dong, G. Zheng, Y. Huang, Y. Chen, L. Li, Z. Xu, N. Li, Z. Liu, Q. Chen, L.-D. Sun and C.-H. Yan, *Science*, 2019, **363**, 265–270.
- Y. Ma, J. Bao, Y. Zhang, Z. Li, X. Zhou, C. Wan, L. Huang, Y. Zhao, G. Han and T. Xue, *Cell*, 2019, **117**, 243–255.
- Z. Li, H. Yuan, W. Yuan, Q. Su and F. Li, *Coord. Chem. Rev.*, 2018, **354**, 155–168.
- Z. Gong, W. Zheng, Y. Gao, P. Huang, D. Tu, R. Li, J. Wei, W. Zhang, Y. Zhang and X. Chen, *Angew. Chem., Int. Ed.*, 2019, **58**, 6943–6947.
- D. Tu, W. Zheng, P. Huang and X. Chen, *Coord. Chem. Rev.*, 2019, **378**, 104–120.
- W. Zheng, P. Huang, Z. Gong, D. Tu, J. Xu, Q. Zou, R. Li, W. You, J.-C. G. Bünzli and X. Chen, *Nat. Commun.*, 2018, **9**, 3462.
- W. Zheng, P. Huang, D. Tu, E. Ma, H. Zhu and X. Chen, *Chem. Soc. Rev.*, 2015, **44**, 1379–1415.
- J. Zhou, Q. Liu, W. Feng, Y. Sun and F. Li, *Chem. Rev.*, 2015, **115**, 395–465.
- F. Wang, R. Deng, J. Wang, Q. Wang, Y. Han, H. Zhu, X. Chen and X. Liu, *Nat. Mater.*, 2011, **10**, 968–973.
- B. Zhou, W. Yang, S. Han, Q. Sun and X. Liu, *Adv. Mater.*, 2015, **27**, 6208–6212.
- S. Fischer, C. Siefe, D. F. Swearer, C. A. McLellan, A. P. Alivisatos and J. A. Dionne, *Angew. Chem., Int. Ed.*, 2020, **59**, 21603–21612.
- H. Dong, L. D. Sun and C. H. Yan, *J. Am. Chem. Soc.*, 2021, **143**, 20546–20561.
- Q. Chen, X. Xie, B. Huang, L. Liang, S. Han, Z. Yi, Y. Wang, Y. Li, D. Fan, L. Huang and X. Liu, *Angew. Chem., Int. Ed.*, 2017, **56**, 7605–7609.
- T. A. Shakhverdov, *Opt. Spectrosc.*, 2003, **95**, 571–580.
- B. Chen and F. Wang, *Acc. Chem. Res.*, 2020, **53**, 358–367.
- Q. Su, H. L. Wei, Y. Liu, C. Chen, M. Guan, S. Wang, Y. Su, H. Wang, Z. Chen and D. Jin, *Nat. Commun.*, 2021, **12**, 4367.
- Q.-C. Sun, H. Mundoor, J. C. Ribot, V. Singh, I. I. Smalyukh and P. Nagpal, *Nano Lett.*, 2014, **14**, 101–106.
- Y. Fan and F. Zhang, *Adv. Opt. Mater.*, 2019, **7**, 1801417.
- Y. Li, S. Zeng and J. Hao, *ACS Nano*, 2019, **13**, 248–259.
- L. Liu, S. Wang, B. Zhao, P. Pei, Y. Fan, X. Li and F. Zhang, *Angew. Chem., Int. Ed.*, 2018, **57**, 7518–7522.
- Y. L. M. Jiang, Z. Xue and S. Zeng, *Theranostics*, 2020, **10**, 6875–6885.
- S. He, J. Song, J. Qu and Z. Cheng, *Chem. Soc. Rev.*, 2018, **47**, 4258–4278.
- J. Xu, A. Gulzar, P. Yang, H. Bi, D. Yang, S. Gai, F. He, J. Lin, B. Xing and D. Jin, *Coord. Chem. Rev.*, 2019, **381**, 104–134.
- B. Zhou, L. Yan, J. Huang, X. Liu, L. Tao and Q. Zhang, *Nat. Photonics*, 2020, **14**, 760–766.
- K. L. Reddy, R. Balaji, A. Kumar and V. Krishnan, *Small*, 2018, **14**, 1801304.
- I. E. Kolesnikov, A. A. Kalinichev, M. A. Kurochkin, E. V. Golyeva, E. Y. Kolesnikov, A. V. Kurochkin, E. Lahderanta and M. D. Mikhailov, *Sci. Rep.*, 2017, **7**, 18002.
- N. Venkatachalam, T. Yamano, E. Hemmer, H. Hyodo, H. Kishimoto and K. Soga, *J. Am. Ceram. Soc.*, 2013, **96**, 2759–2765.
- Y. Zhong, Z. Ma, S. Zhu, J. Yue, M. Zhang, A. L. Antaris, J. Yuan, R. Cui, H. Wan, Y. Zhou, W. Wang, N. F. Huang, J. Luo, Z. Hu and H. Dai, *Nat. Commun.*, 2017, **8**, 737.
- H. Zhang, Y. Fan, P. Pei, C. Sun, L. Lu and F. Zhang, *Angew. Chem., Int. Ed.*, 2019, **58**, 10153–10157.

- 36 H. Li, X. Wang, T. Y. Ohulchanskyy and G. Chen, *Adv. Mater.*, 2021, **33**, 2000678.
- 37 Y. Xu, X. Gong, Y. Chen, M. Huang, Z. Luo and Y. Huang, *J. Cryst. Growth*, 2003, **252**, 241–245.
- 38 S. Wang, B. Shen, H. L. Wei, Z. Liu, Z. Chen, Y. Zhang, Y. Su, J. Z. Zhang, H. Wang and Q. Su, *Nanoscale*, 2019, **11**, 10220–10228.
- 39 D. Hudry, A. D. Backer, R. Popescu, D. Busko, I. A. Howard, S. Bals, Y. Zhang, A. Pedrazo-Tardajos, S. V. Aert, D. Gerthsen, T. Altantzis and B. S. Richards, *Small*, 2021, **17**, 2104441.
- 40 D. Hudry, R. Popescu, D. Busko, M. Diaz-Lopez, M. Abeykoon, P. Bordet, D. Gerthsen, I. A. Howard and B. S. Richards, *J. Mater. Chem. C*, 2019, **7**, 1164–1172.
- 41 D. Hudry, I. A. Howard, R. Popescu, D. Gerthsen and B. S. Richards, *Adv. Mater.*, 2019, **31**, 1900623.
- 42 D. Hudry, D. Busko, R. Popescu, D. Gerthsen, I. A. Howard and B. S. Richards, *J. Mater. Chem. C*, 2019, **7**, 7371–7377.
- 43 L. Liu, X. Li, Y. Fan, C. Wang, A. M. El-Toni, M. S. Alhoshan, D. Zhao and F. Zhang, *Chem. Mater.*, 2019, **31**, 5608–5615.
- 44 P. U. Bastian, S. Nacak, V. Roddatis and M. U. Kumke, *J. Phys. Chem. C*, 2020, **124**, 11229–11238.
- 45 W.-P. Qin, C.-Y. Cao, L.-L. Wang, J.-S. Zhang, D.-S. Zhang, K.-Z. Zheng, Y. Wang, G.-D. Wei, G.-F. Wang, P.-F. Zhu and R.-J. Kim, *Opt. Lett.*, 2008, **33**, 2167–2169.
- 46 K. Zheng, W. Qin, C. Cao, D. Zhao and L. Wang, *J. Phys. Chem. Lett.*, 2015, **6**, 556–560.
- 47 W.-P. Qin, Z.-Y. Liu, C.-N. Sin, C.-F. Wu, G.-S. Qin, Z. Chen and K.-Z. Zheng, *Light: Sci. Appl.*, 2014, **3**, e193.
- 48 M. J. Weber, *Phys. Rev. B: Solid State*, 1971, **4**, 3153–3159.
- 49 K. Bartosiewicz, V. Babin, K. Kamada, A. Yoshikawa and M. Nikl, *J. Lumin.*, 2015, **166**, 177–122.
- 50 R. Balda, J. I. Peña, M. A. Arriandiaga and J. Fernández, *Opt. Express*, 2010, **18**, 13842–13850.
- 51 W. Wei, G. Chen, A. Baev, G. S. He, W. Shao, J. Damasco and P. N. Prasad, *J. Am. Chem. Soc.*, 2016, **138**, 15130–15133.
- 52 W. Shao, G. Chen, A. Kuzmin, H. L. Kutscher, A. Pliss, T. Y. Ohulchanskyy and P. N. Prasad, *J. Am. Chem. Soc.*, 2016, **138**, 16192–16195.
- 53 W. Holzer, M. Mauerer, A. Penzkofer, R.-M. Szeimies, C. Abels, M. Landthaler and W. Bäuml, *J. Photochem. Photobiol., B*, 1998, **47**, 155–164.
- 54 V. Saxena, M. Sadoqi and J. Shao, *J. Photochem. Photobiol., B*, 2004, **74**, 29–38.
- 55 V. Muhr, C. Würth, M. Kraft, M. Buchner, A. J. Baeumner, U. Resch-Genger and T. Hirsch, *Anal. Chem.*, 2017, **89**, 4868–4874.
- 56 D. J. Garfield, N. J. Borys, S. M. Hamed, N. A. Torquato, C. A. Tajon, B. Tian, B. Shevitski, E. S. Barnard, Y. D. Suh, S. Aloni, J. B. Neaton, E. M. Chan, B. E. Cohen and P. J. Schuck, *Nat. Photonics*, 2018, **12**, 402–407.
- 57 D. Dolmans, D. Fukumura and R. Jain, *Nat. Rev. Cancer*, 2003, **3**, 380–387.
- 58 W. S. Kuo, Y. T. Chang, K. C. Cho, K. C. Chiu, C. H. Lien, C. S. Yeh and S. J. Chen, *Biomaterials*, 2012, **33**, 3270–3278.
- 59 Z. Hou, Y. Zhang, K. Deng, Y. Chen, X. Li, X. Deng, Z. Cheng, H. Lian, C. Li and J. Lin, *ACS Nano*, 2015, **9**, 2584–2599.