

Supplementary Information for

Three-primary-color up-conversion luminescence from single tri-
sensitized NaYF₄ nanocrystals

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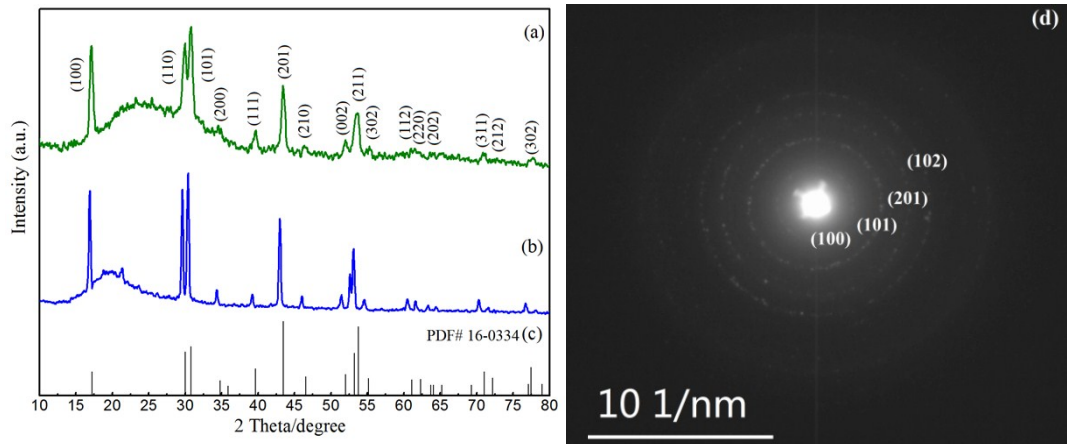


Figure S1. XRD patterns of (a) NaYF₄: Nd, Yb, Tm and (b) NaYF₄: Nd, Yb, Tm@ NaYF₄: Nd samples, which suggest the crystal structure of β -NaYF₄. (c) stander cards of β -NaYF₄ (PDF# 16-0334). (d) Electron diffraction of as-prepared C-S1-S2-S3-S4 UC nanoparticles indicates the β -NaYF₄ crystals.

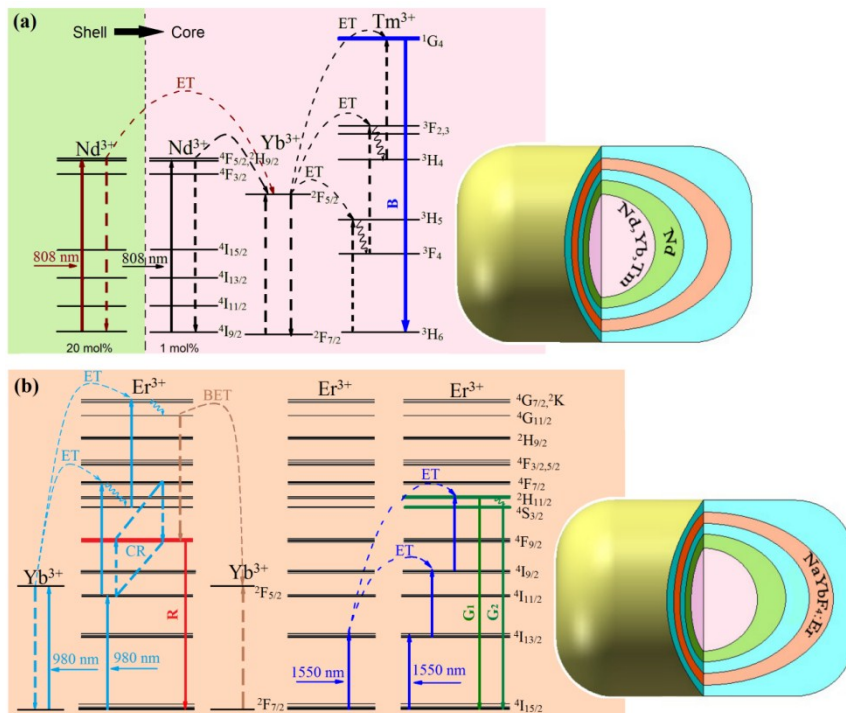


Figure S2. UC luminescence mechanisms of NaYF₄-based core-shell nanoparticles. (a) The C and S1 layers exhibit blue emission ($^1G_4 \rightarrow ^3H_6$) via energy transfer (ET) processes of Nd³⁺ \rightarrow Yb³⁺ \rightarrow Tm³⁺ under 808 nm excitation. (b) The S3 layer of NaYbF₄: Er³⁺ presents individual red (marked as R), and green (marked as G1, G2) luminescence. Cross relaxation (CR) between Er³⁺ ions and back energy transfer (BET) from Er³⁺ \rightarrow Yb³⁺ ions lead to red emission excited at 980 nm. While at 1550 nm excitation, continuous absorption of three photons or ET processes between Er³⁺ ions produce green emission.

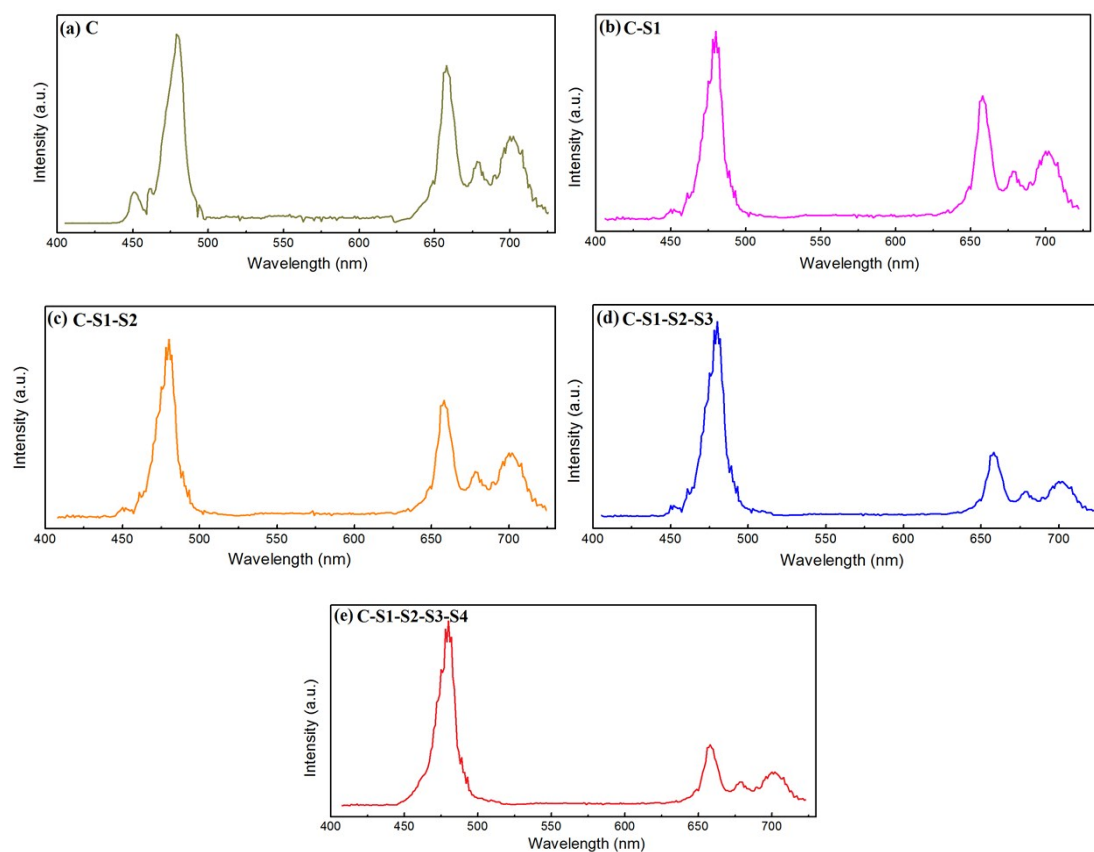
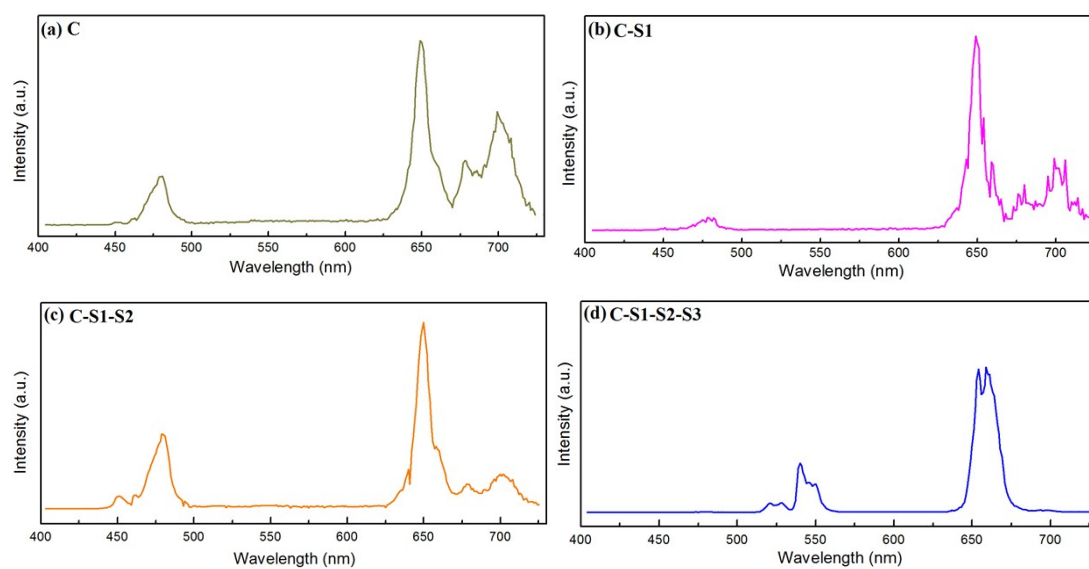


Figure S3. UC luminescence spectra of (a) C, (b) C-S1, (c) C-S1-S2, (d) C-S1-S2-S3, (e) C-S1-S2-S3-S4 UC nanoparticles under 808 nm excitation. The blue emission is enhanced with the shell coating at excitation of 808 nm.



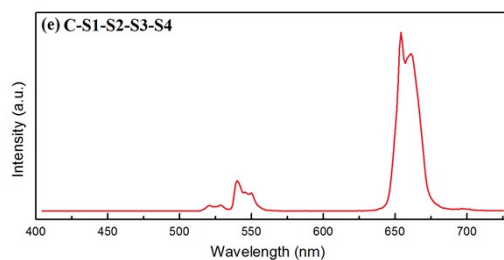


Figure S4. UC luminescence spectra of (a) C, (b) C-S1, (c) C-S1-S2, (d) C-S1-S2-S3, (e) C-S1-S2-S3-S4 UC nanoparticles under 980 nm excitation. The blue emission is suppressed with the shell coating at excitation of 980 nm.

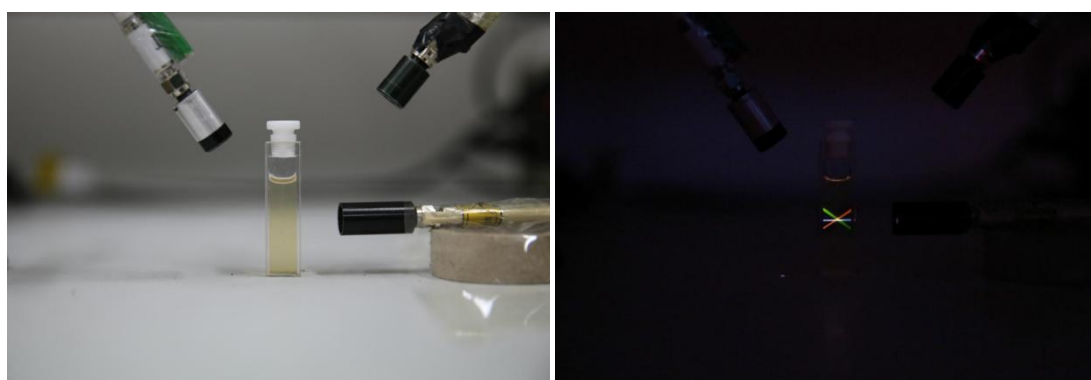


Figure S5. Photographs of three-primary-color luminescence from as-prepared multilayer β -NaYF₄ nanoparticles. (Canon EOS 5D Mark III, Tv=1:10, Len: EF24-70mm f/2.8L II USM, Av= 2.8)

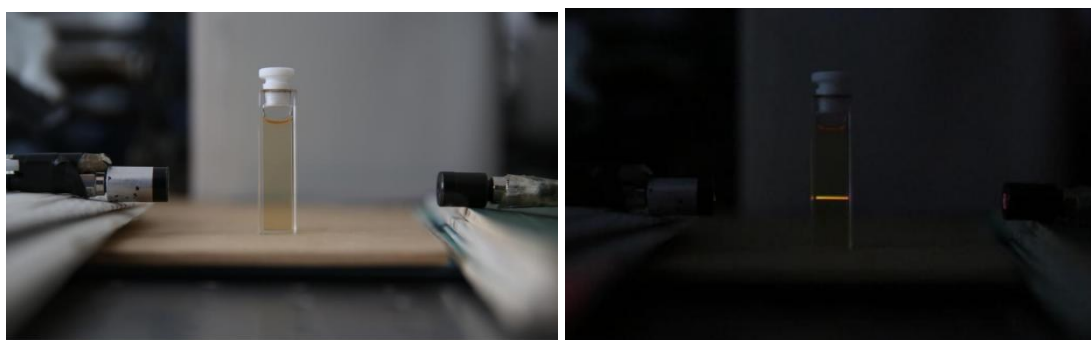


Figure S6. An example of camera images that demonstrates tunable UC luminescence measurement of multilayer β -NaYF₄ nanoparticles. (Canon EOS 5D Mark III, Tv=1:10, Len: EF24-70mm f/2.8L II USM, Av= 2.8).



Video.mp4