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Supporting Information

Regulation of multifunctional mesoporous core-shell nanoparticles with luminescence and magnetic properties for biomedical applications

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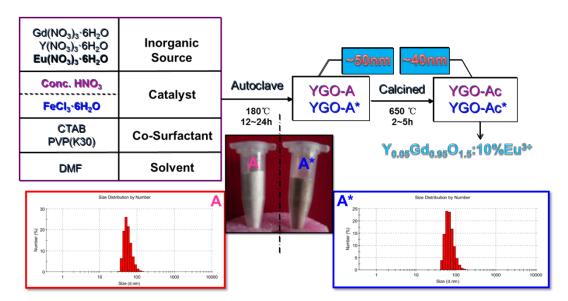


Figure S1. Flow chart of solvothermal synthetic procedures for $(Y,Gd)_2O_3$:Eu³⁺ NPs. The inset at the bottom: DLS size distribution (in ethanol) of YGO-A NPs (left) and YGO-A* NPs (right).

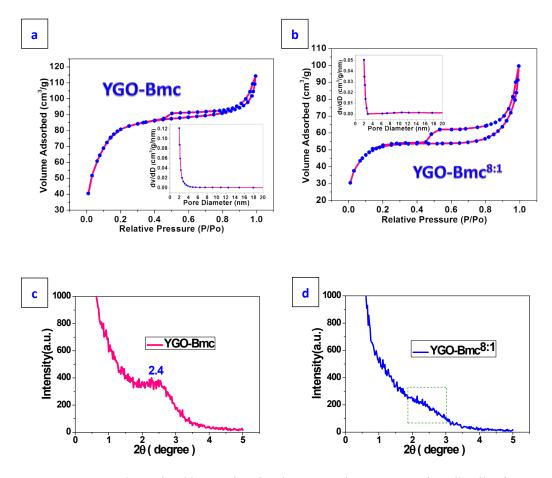


Figure S2. N₂ adsorption/desorption isotherms and mesopore size distribution curves (the inset) of (a) YGO-Bmc NPs, (b) YGO-Bmc^{8:1} NPs; low-angle XRD patterns of (c) YGO-Bmc NPs, (d) YGO-Bmc^{8:1} NPs.

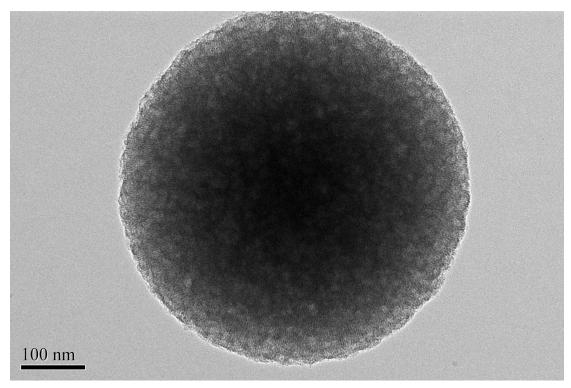


Figure S3. TEM image of $mSiO_2$.

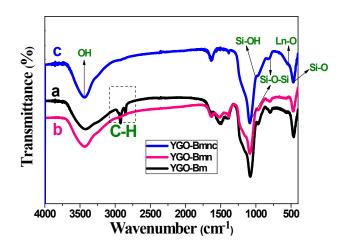


Figure S4. The FT-IR spectra of (a) YGO-Bm NPs, (b) YGO-Bmn NPs, (c) YGO-Bmnc NPs. Contrary to YGO-Bm NPs, the bands in the region 2800~3000 cm⁻¹ which attributed to the vibrations of -CH₃, -CH₂ of CTAB templates disappeared for YGO-Bmn NPs, suggesting that the CTAB templates are eradicated completely after NH₄NO₃/EtOH extraction. For YGO-Bmnc NPs, the new characteristic vibration of Ln-O bond in 543 cm⁻¹ turns up, demonstrating that the amorphous Ln(OH)₃ completely dehydrated and transformed into Ln₂O₃ after calcination.

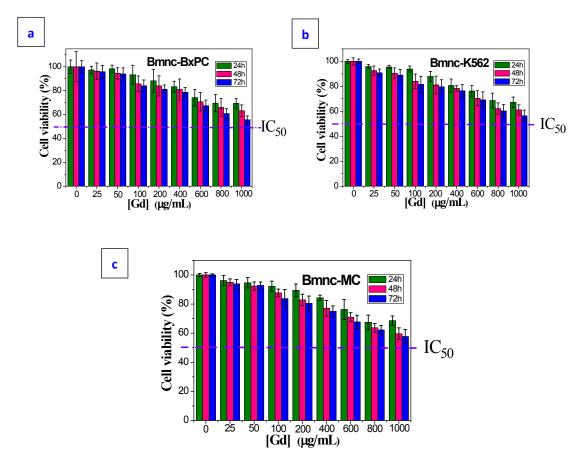


Figure S5. In vitro cytotoxicity of YGO-Bmnc NPs against BxPC-3, K562 and MC cells after 24 h, 48 h and 72h incubation.