

## Supporting Information

# Reusable and pH-Stable Luminescent Sensors for Highly Selective Detection of Phosphate

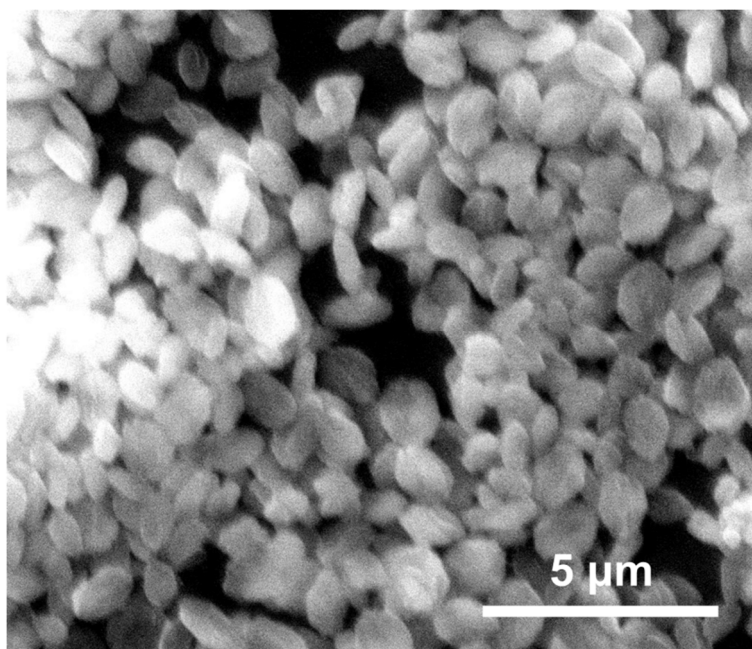
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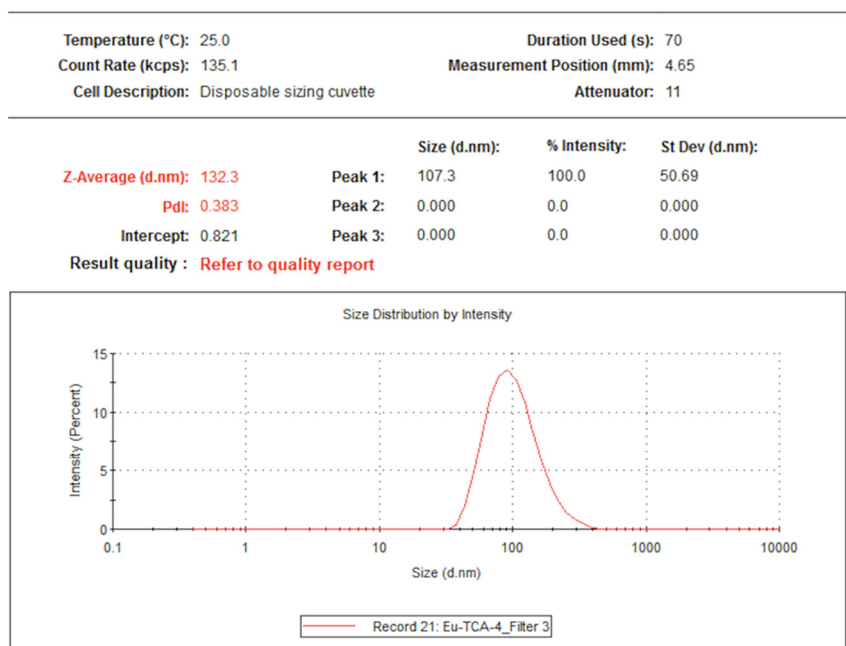
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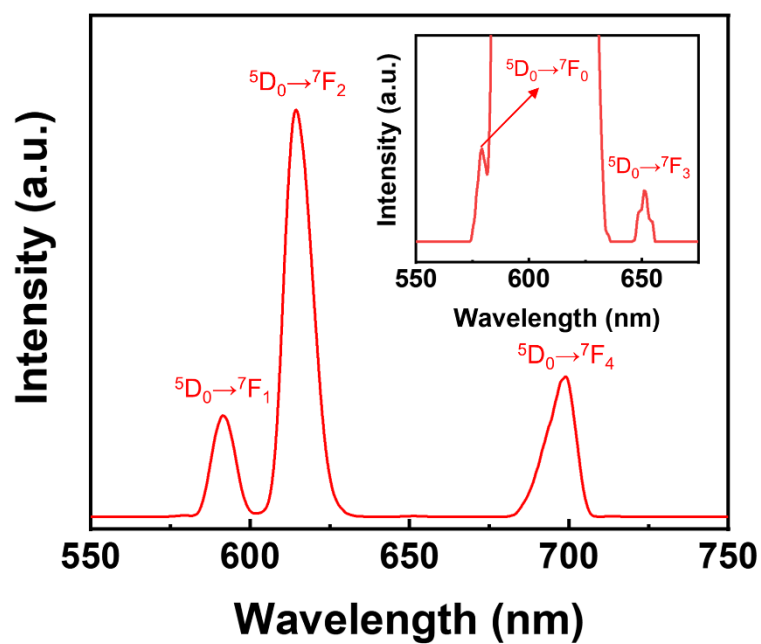
**Figure S1.** SEM image of Eu-TCA microcrystals.



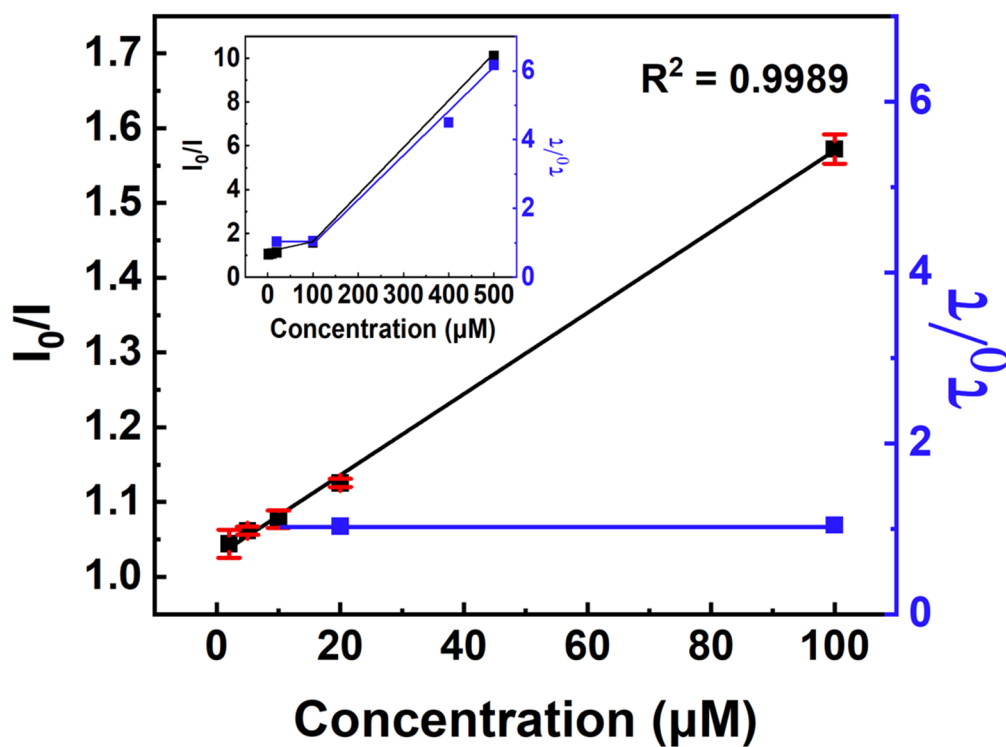
**Figure S2.** Dynamic light scattering analysis of Eu-TCA.

### Synthesis of Eu-TCA microcrystals

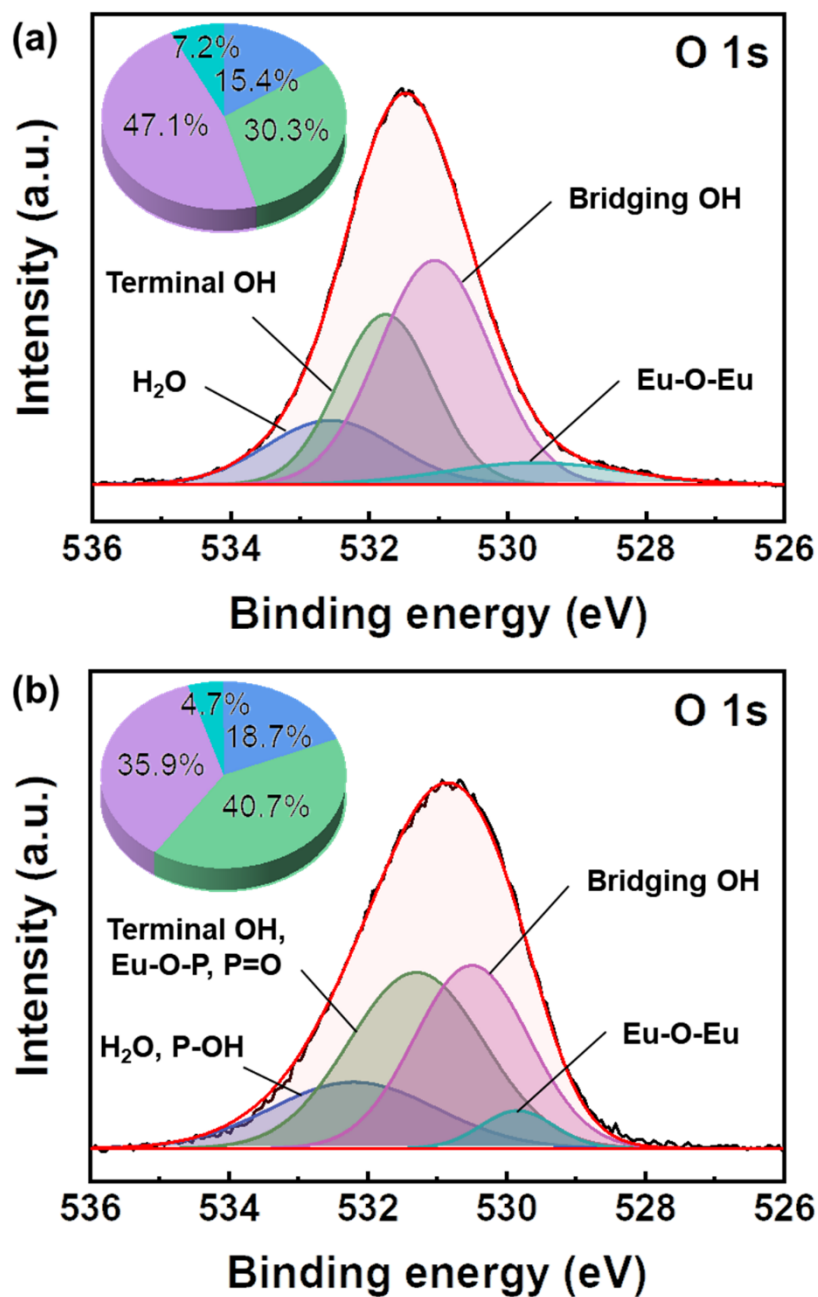
EuCl<sub>3</sub> (235 mg, 0.64 mmol, 5.3 eq.), TCA ligand (60 mg, 0.12 mmol, 1 eq.), and polyvinylpyrrolidone (533 mg) were reacted under solvothermal conditions in a mixed solvent of dimethylformamide (32 mL) and ethanol (19 mL). The reaction was conducted at 150°C for 12 h. After the reaction, the reaction mixture was cooled to room temperature over the next 6 h, and colloidal particles were precipitated by centrifugation. Three cycles of washing with dry dimethylformamide and centrifugation were followed by a final wash with ethanol. The sample was dried in an oven at 70°C for 3 h.



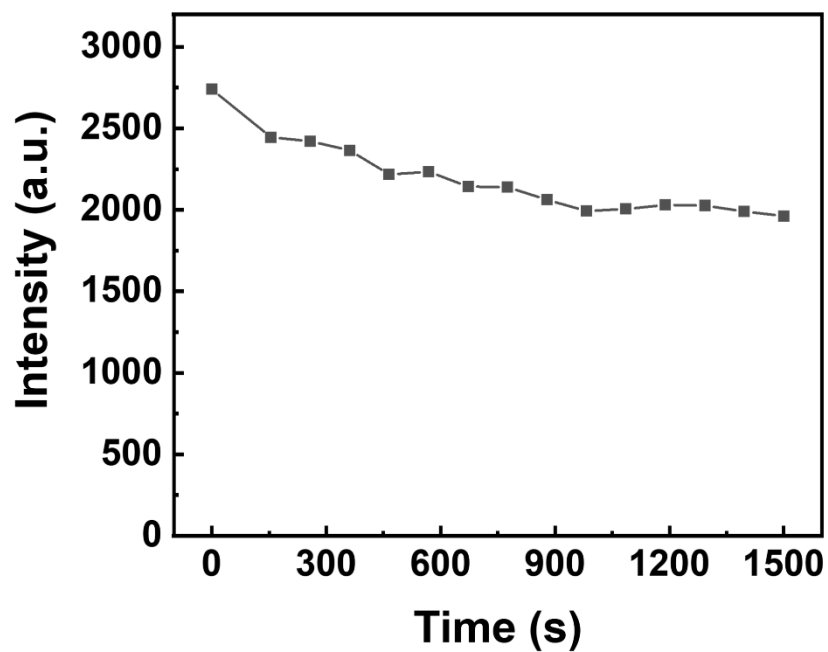
**Figure S3.** Emission spectrum of Eu-TCA dispersion ( $\lambda_{\text{ex}} = 260$  nm). The inset shows enlarged emission spectrum.



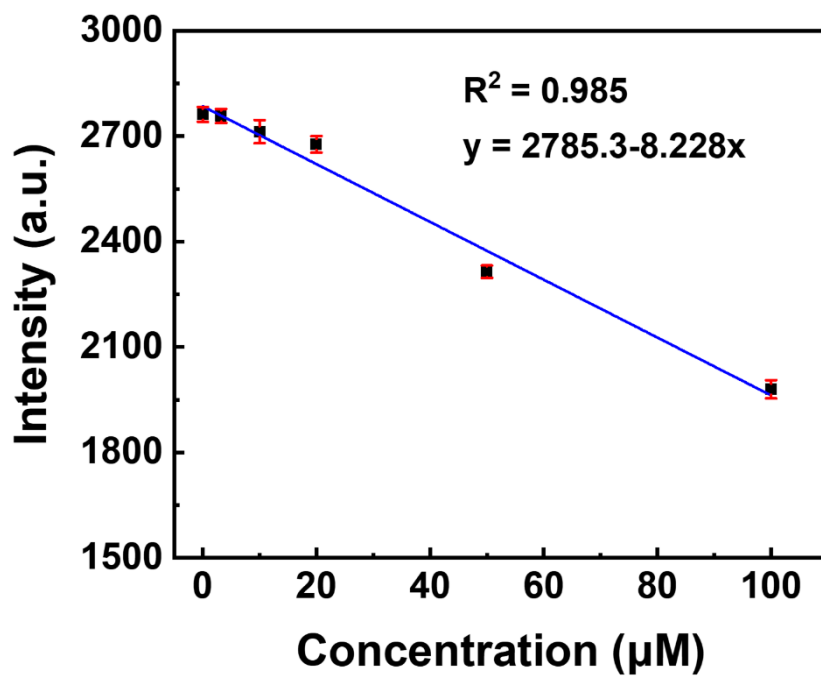
**Figure S4.** *Stern-Volmer* plot for the luminescence quenching of Eu-TCA dispersion upon addition of different concentrations of phosphate. Inset shows the *Stern-Volmer* plot in the concentration range of 2–500  $\mu\text{M}$ .



**Figure S5.** Experimental fit performed on O 1s XPS spectra before (a) and after (b) incubation with phosphate.



**Figure S6.** Time-dependent luminescence intensity of the Eu-TCA/GMF upon the addition of 100 μM of phosphate.



**Figure S7.** The luminescence intensity of Eu-TCA/GMF under various phosphate concentrations and their linear fit curve for the estimation of LOD.

**Table S1.** Comparison of various analytical methods for phosphate detection.

Analytical methods	Sensing element/structure	Linear range ( $\mu\text{M}$ )	LOD ( $\mu\text{M}$ )	Reusable <sup>1</sup>	pH range <sup>2</sup>	Ref.
Potentiometry	Mo-based electrode	10–10 <sup>5</sup>	1.9	O	pH-unstable <sup>3</sup>	S1
	W-based electrode	1–10 <sup>5</sup>	0.4	O	pH-unstable <sup>3</sup>	S2
	Polymeric membrane electrode	32–10 <sup>5</sup>	10	O	pH-unstable <sup>3</sup>	S3
Voltammetry	Screen-printed electrodes modified with carbon black nanoparticles	0.5–100	0.1	O	pH-unstable <sup>3</sup>	S4
I-V measurement	Ag/graphene composite-based field-effect transistor	5–6000	1.2	O	–	S5
Colorimetric probe	Au nanoparticles/Eu <sup>3+</sup>	0.5–30	0.076	X	–	S6
	Functionalized Au nanoparticles	80–200	120	X	7–8	S7
Luminescent probe	Eu@BUC-14	5–150	0.88	X	4–8	S8
	UiO-66-NH <sub>2</sub> MOF <sup>4</sup>	5–150	1.25	X	–	S9
	Eu-triazole MOF <sup>4</sup>	3–30	6.62	X	–	S10
	Tb-MOF <sup>4</sup>	40–400	35	X	–	S11
	Eu-based nanospheres	2–100	0.83	X	5–9	S12
	Eu-TCA/GMF	3–500	1.52	O	3–10	This work

<sup>1</sup> The term “reusable” means the sensing material can be easily reused without complex processes such as centrifugation or drying. <sup>2</sup> pH range that does not affect the sensor performance. <sup>3</sup> Needs pH compensation model. <sup>4</sup> MOF: metal-organic framework.

## References

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