

Fluorescein-based ionic liquid sensor for label-free detection of serum albumins

Supporting Information

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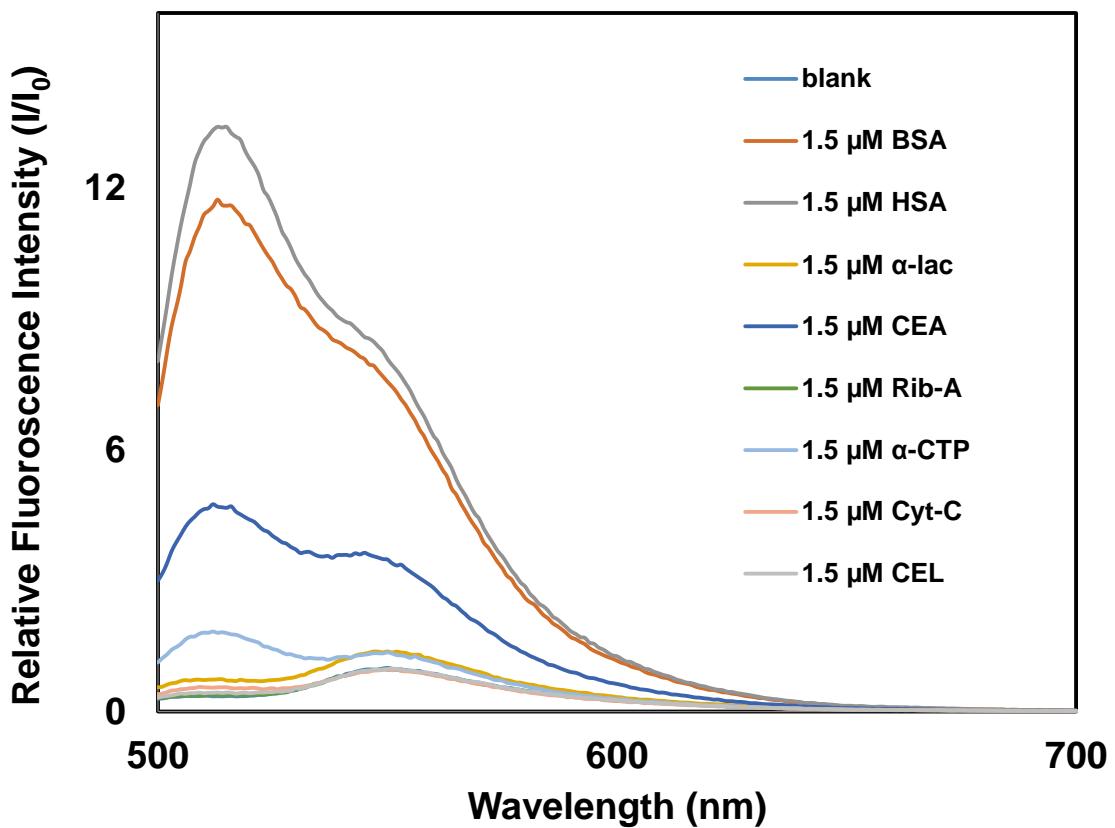
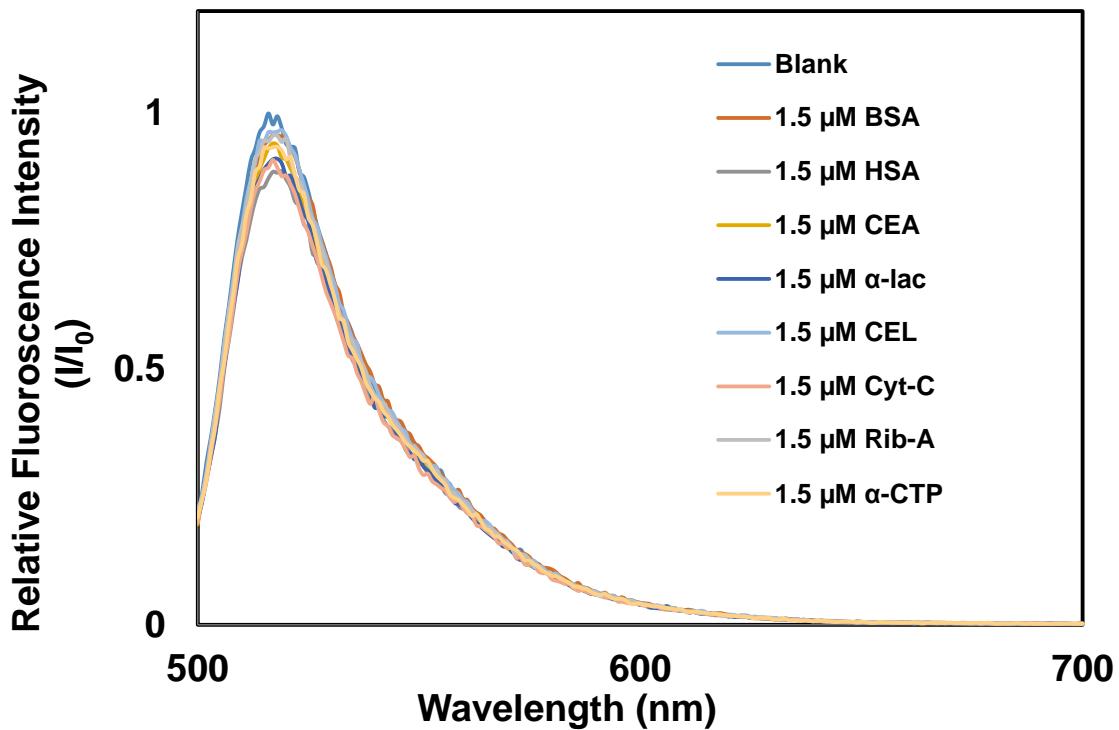
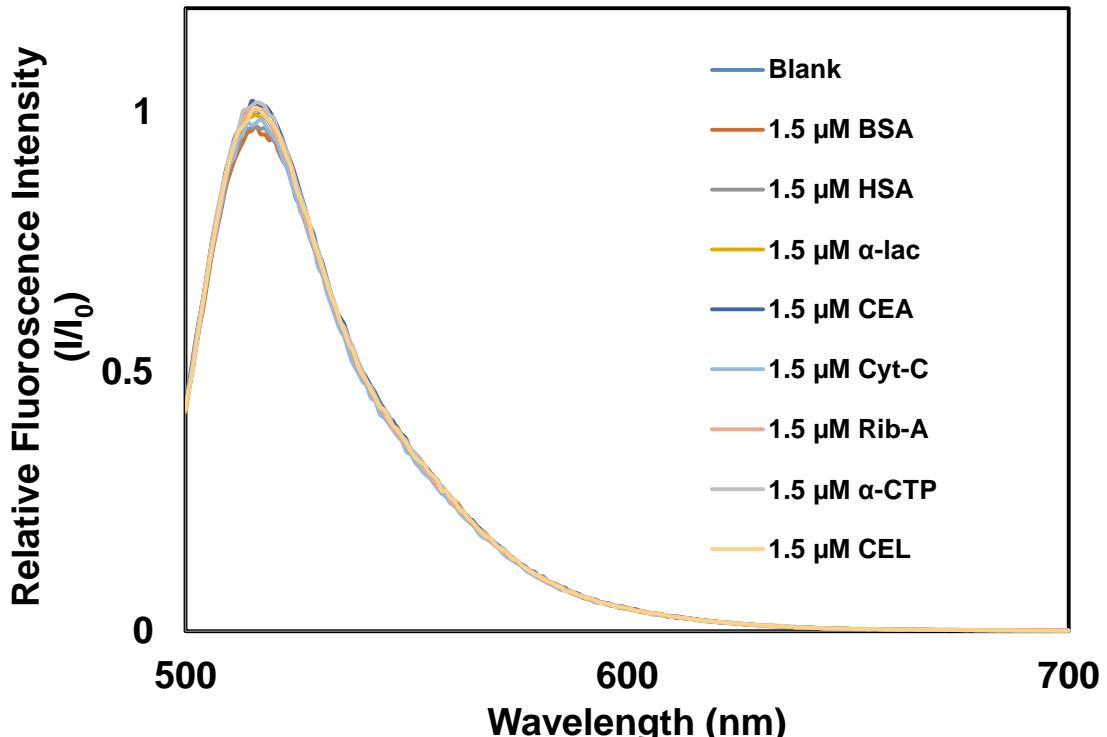


Fig. S1. Fluorescence emission spectra ($\lambda_{\text{ex}} = 490 \text{ nm}$) of $40 \mu\text{M} [\text{P}_{66614}]_2[\text{FL}]$ nanodroplets in the presence of same concentration ($1.5 \mu\text{M}$) of different albumins and non-albumins

a



b



C

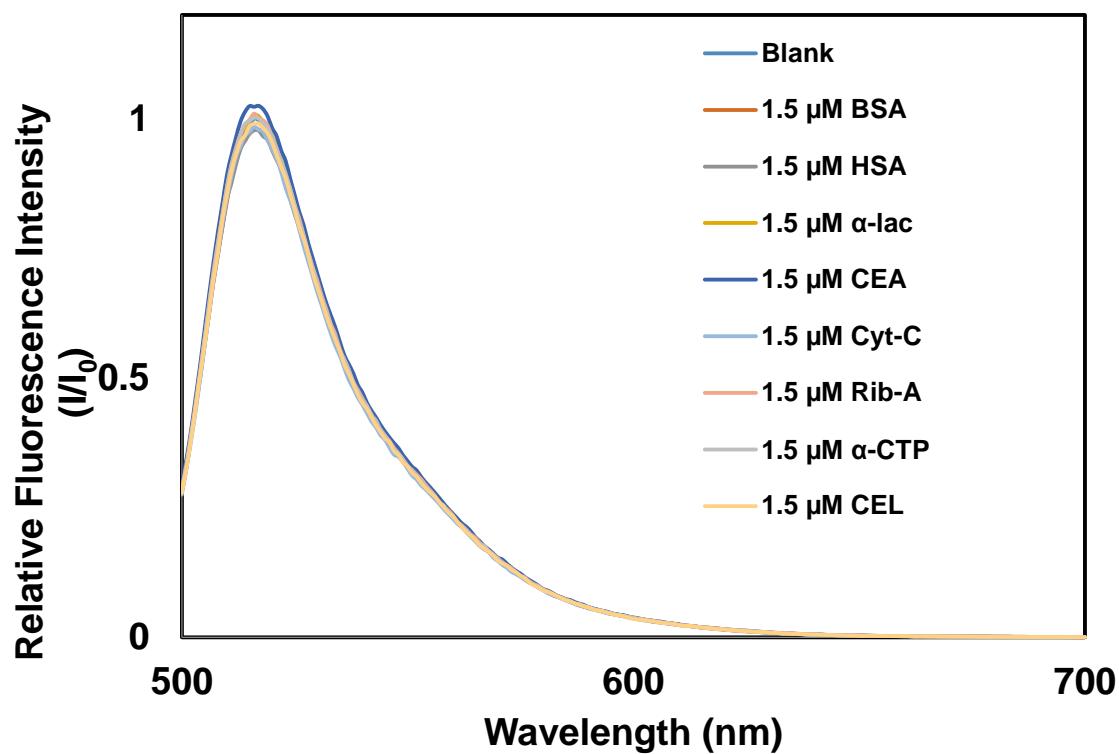
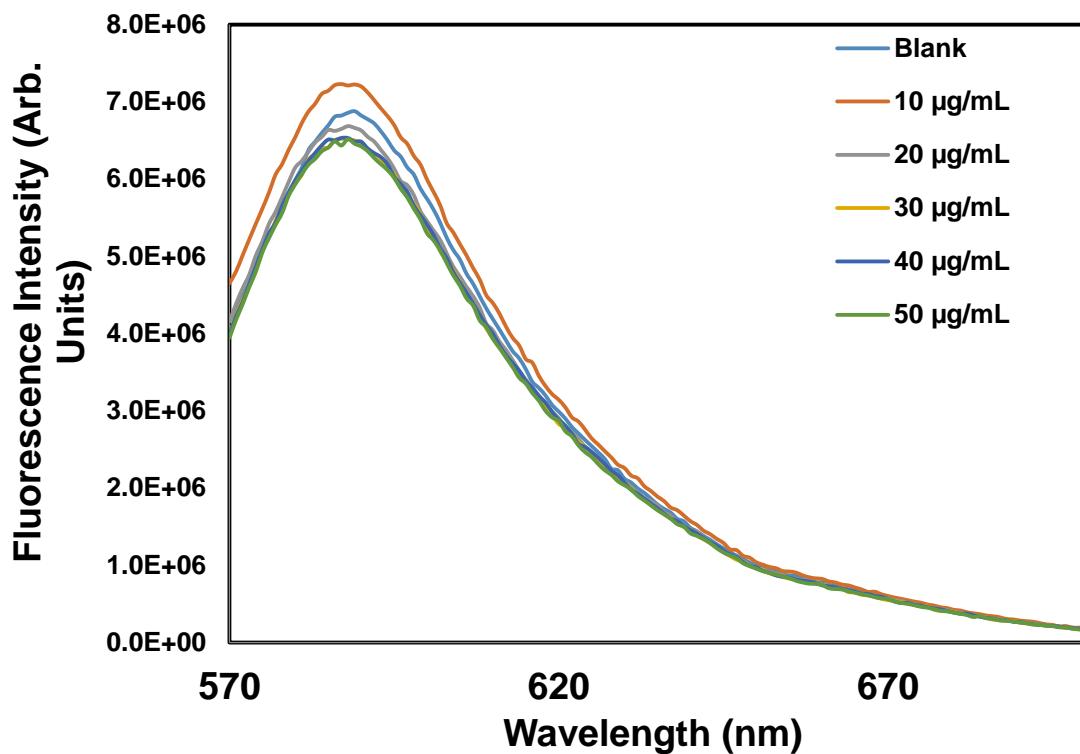
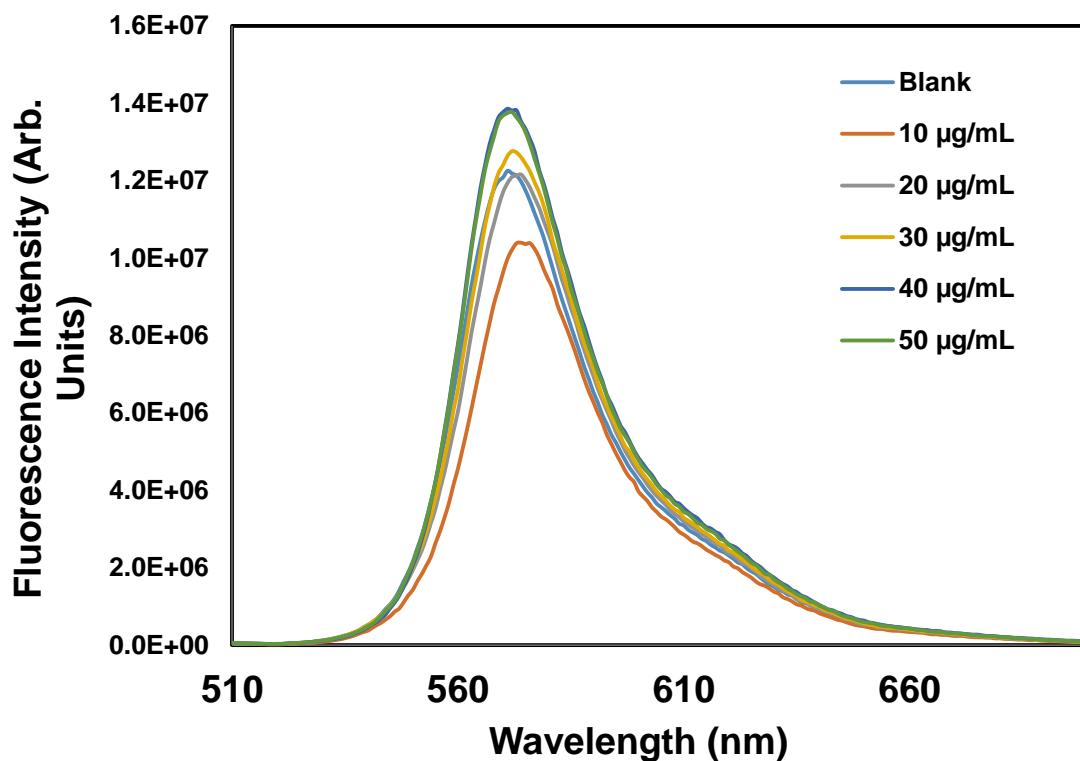


Fig. S2. Fluorescence emission spectra ($\lambda_{\text{ex}} = 490$ nm) of 40 μ M (A) Na_2FL , (B) $[\text{TPP}]_2[\text{FL}]$ and (C) $[4\text{NB}]_2[\text{FL}]$ with eight different proteins at the concentration of 1.5 μ M

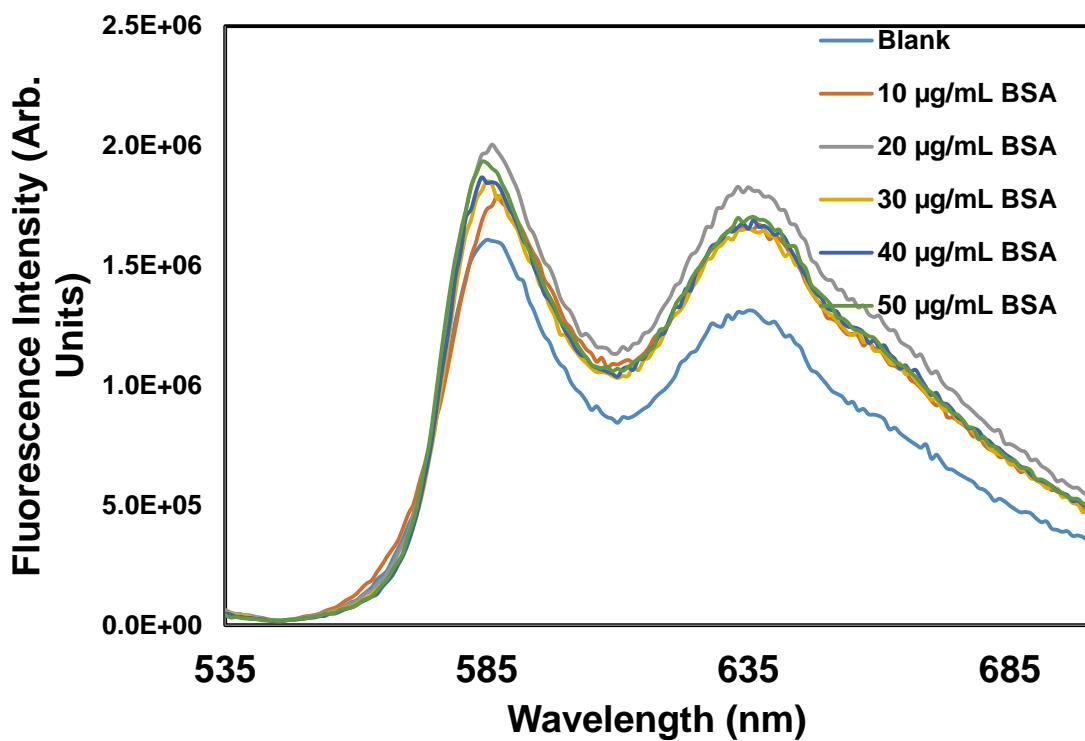
a



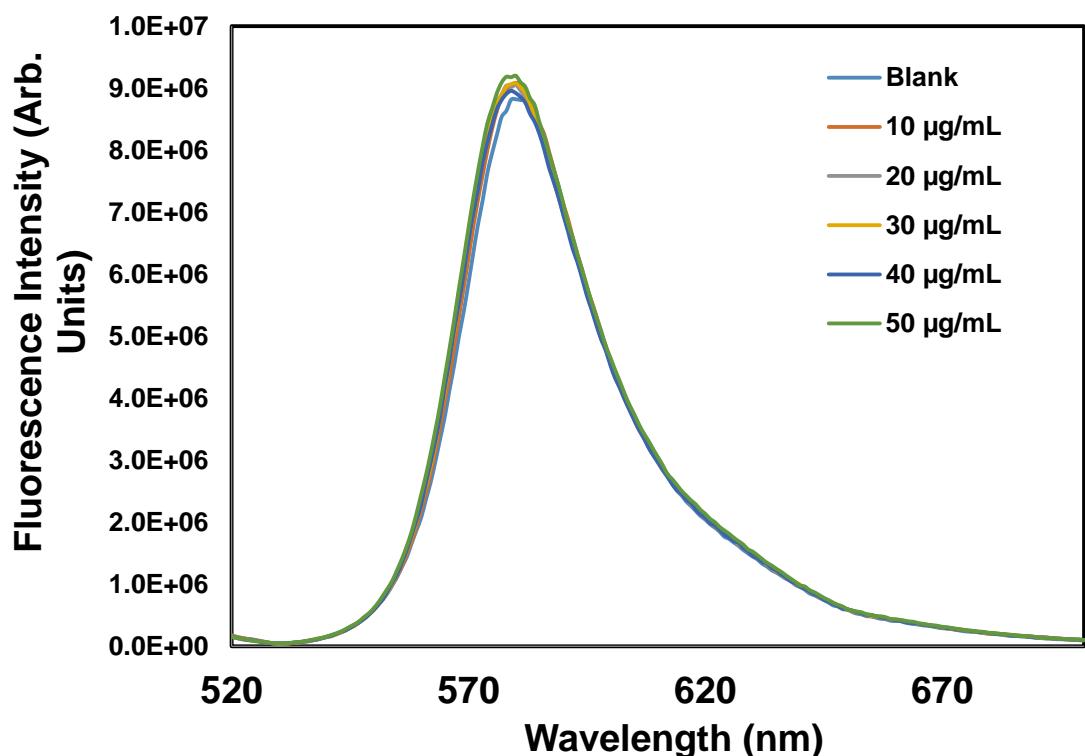
b



c



d



e

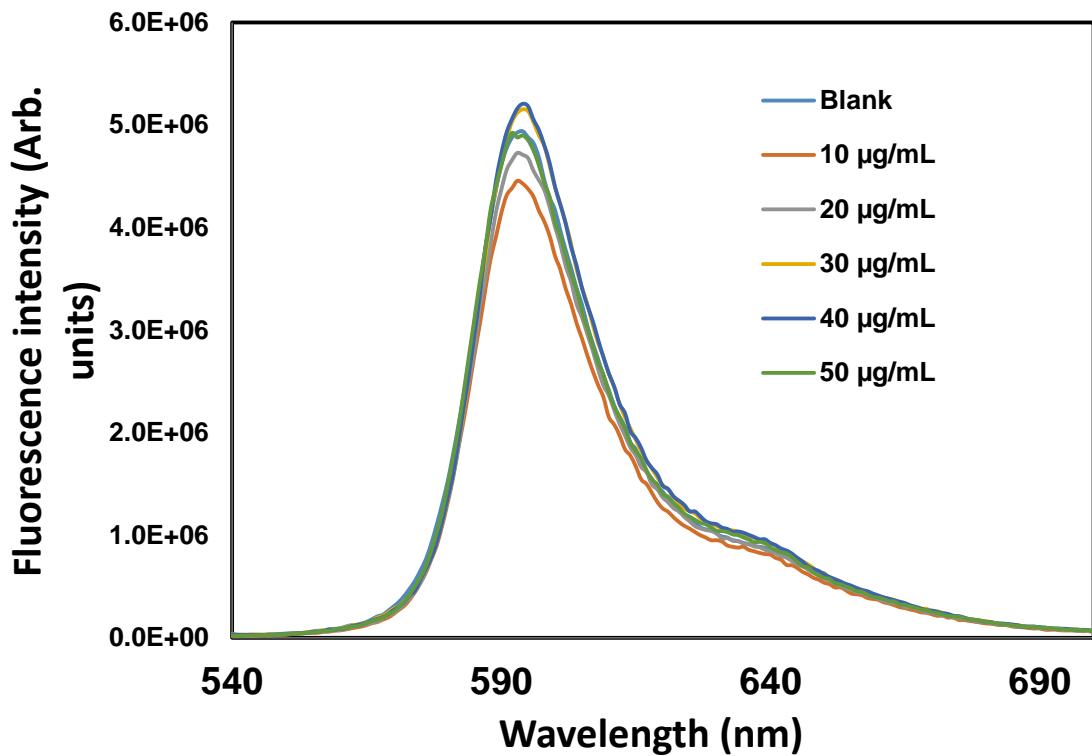


Fig. S3. Fluorescence emission spectra ($\lambda_{\text{ex}} = 490$ nm) of (a) Eosin B, (b) Eosin Y, (c) Phloxine B (d) Erythrosin B and, (e) Rose Bengal nanoparticles dispersed in different concentrations of BSA

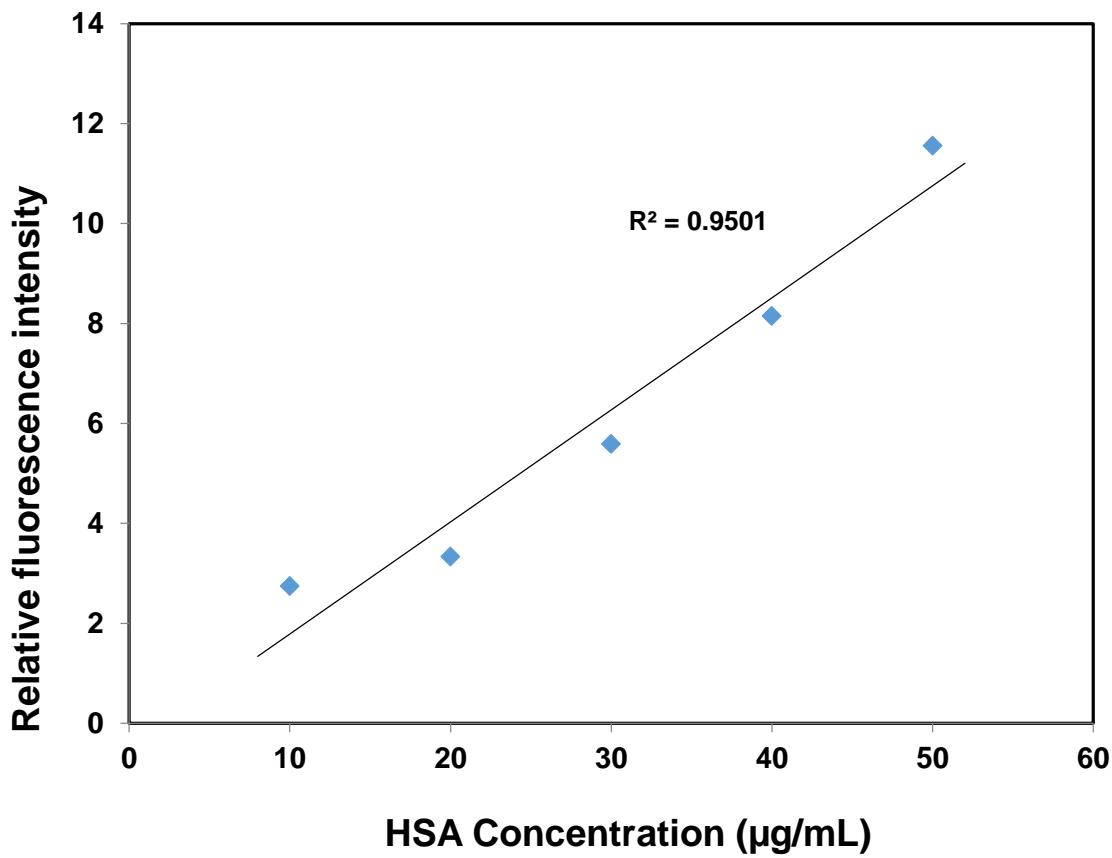


Fig S4. Relationship between relative fluorescence intensity at 512 nm and HSA concentration in human serum. Human serum samples were diluted for 1000 times before analysis. (concentration of $[P_{66614}]_2[FL]$ - 24 µM)

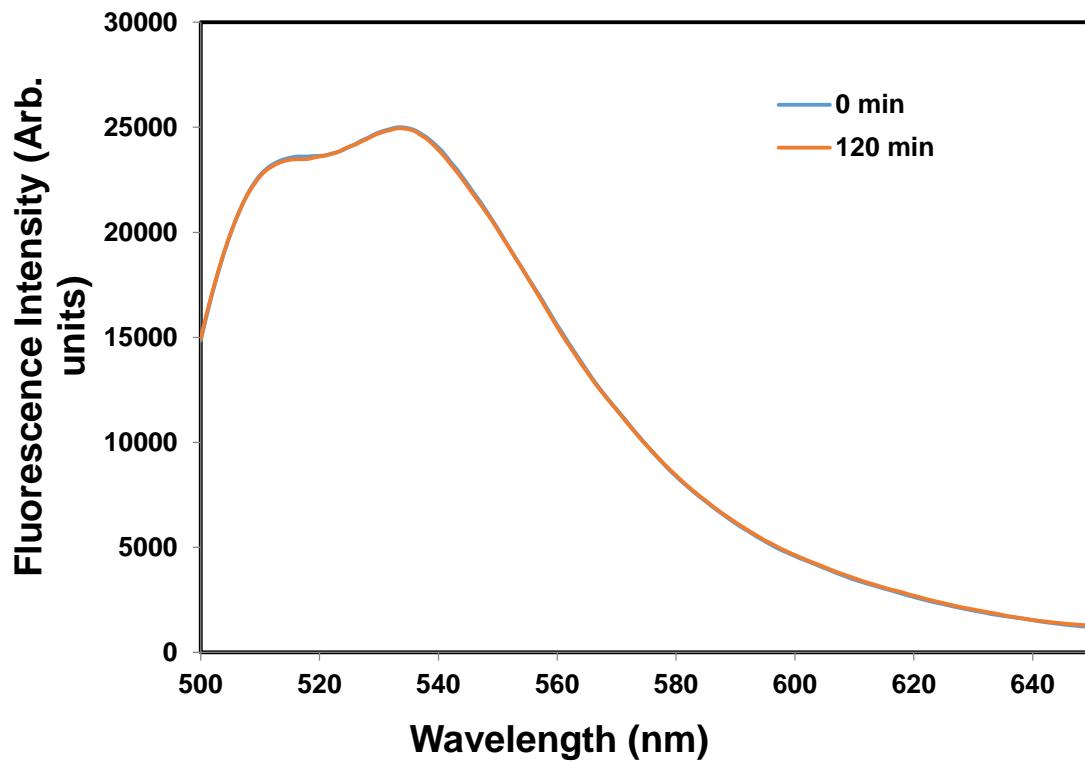


Fig. S5. The fluorescence emission spectra ($\lambda_{\text{ex}} = 490 \text{ nm}$) of $24 \mu\text{M}$ dispersions of $[\text{P}_{66614}]_2[\text{FL}]$ at 0 and 120 minutes

Table S1. Aggregate to monomer peak ratio $A_{\text{ag}(530)}/A_{\text{m}(490)}$ derived from absorbance spectra

[P ₆₆₆₁₄] ₂ [FL] Concentration(μM)	$A_{\text{ag}(530)}/A_{\text{m}(490)}$ ratio
40	1.9
32	1.9
24	1.9
16	1.6
08	1.1

A_{ag} – Absorbance of the aggregate peak, A_{m} – Absorbance of the monomer peak