Deciphering the Binding Behaviours of BSA Using Ionic AIE-active Fluorescent Probes

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Figure S1. Conformational structure of BSA in its native state.



Figure S2. (A) FL spectra of BSA (1 μ M) excited at 280 nm in the presence of different concentrations of M3; (B) Plot of lg [(F_0 -F)/F] vs. lg [M3]. F_0 and F: the peak FL intensity of BSA without the probe and with different concentrations of the probe.



Figure S3. (A) FL spectra of BSA+M2 system in the presence of different concentrations of myristic acid (0~10 μ M). (B) Plots of the corresponding fluorescent intensities. Excitation wavelength (λ_{ex}) : 330 nm. Concentration of M2: 1 μ M; Concentration of BSA: 1 μ M.



Figure S4. (A) FL spectra of BSA+M**3** with different concentrations ($0 \sim 10 \mu$ M) of myristic acid; (B) FL spectra of M**3** with different concentrations ($0 \sim 10 \mu$ M) of myristic acid; (C) Plots of the corresponding fluorescent intensities. λ_{ex} : 395 nm. Concentration of M**3**: 1 μ M; Concentration of BSA: 1 μ M.



Figure S5. FL spectra of BSA+M2 system in PBS buffer (pH=7.4) at different temperatures (heating from 25 °C to 75 °C, and then cooling from 75 °C to 25 °C). λ_{ex} : 330 nm. Concentration of M2: 5 μ M; Concentration of BSA: 1 μ M.



Figure S6. FL spectra of BSA+M**3** system in PBS buffer (pH=7.4) at different temperatures (heating from 25 °C to 75 °C, and then cooling from 75 °C to 25 °C). λ_{ex} : 395 nm. Concentration of M**3**: 5 μ M; Concentration of BSA: 1 μ M.



Figure S7. FL spectra of BSA+M2 system in PBS buffer (pH=7.4) at different temperatures (heating from 25 °C to 50 °C and then cooling from 50 °C to 25 °C). λ_{ex} : 330 nm. Concentration of M2: 5 μ M; Concentration of BSA: 1 μ M.



Figure S8. FL spectra of BSA+M**3** system in PBS buffer (pH=7.4) at different temperatures (heating from 25 °C to 50 °C, and then cooling from 50 °C to 25 °C). λ_{ex} : 395 nm. Concentration of M**3**: 5 μ M; Concentration of BSA: 1 μ M.



Figure S9. FL spectra of BSA+M2 system in the presence of different concentrations of GndHCl. λ_{ex} : 330 nm; Concentration of M2: 1 μ M; Concentration of BSA: 1 μ M.



Figure S10. FL spectra of BSA+M3 system in the presence of different concentrations of GndHCl. λ_{ex} : 395 nm; Concentration of M3: 1 μ M; Concentration of BSA: 1 μ M.



Figure S11. FL spectra of BSA+M2 system in the presence of different concentrations of urea. λ_{ex} : 330 nm; Concentration of M2: 1 μ M; Concentration of BSA: 1 μ M.



Figure S12. FL spectra of BSA+M**3** system in the presence of different concentrations of urea. λ_{ex} : 395 nm; Concentration of M**3**: 1 μ M; Concentration of BSA: 1 μ M.



Figure S13. FL spectra of M2 in the presence (A) or absence (B) of BSA in buffer with different pH values (2.0~7.4). λ_{ex} : 330 nm; Concentration of M2: 1 μ M; Concentration of BSA: 1 μ M.



Figure S14. FL spectra of M**3** in the presence (A) or absence (B) of BSA in buffer with different pH values (2.0~7.4). λ_{ex} : 395 nm; Concentration of M**3**: 1 μ M; Concentration of BSA: 1 μ M.