Supplementary Material for Organic & Biomolecular Chemistry

Recognition and sensing of solvent exposed protein surfaces using anthracene derived receptors

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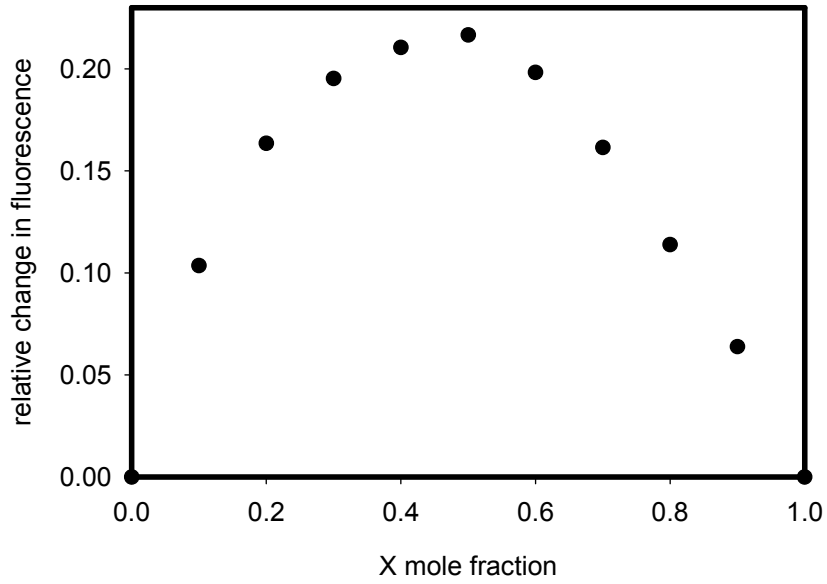
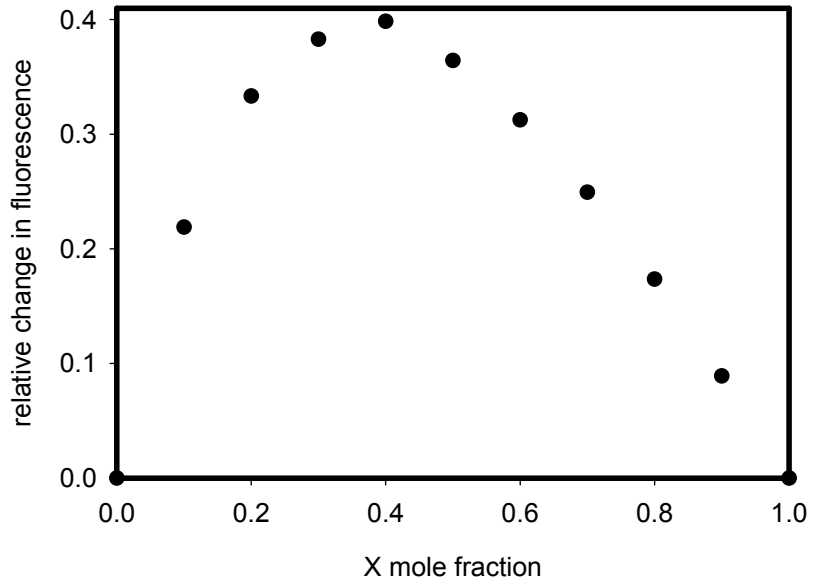
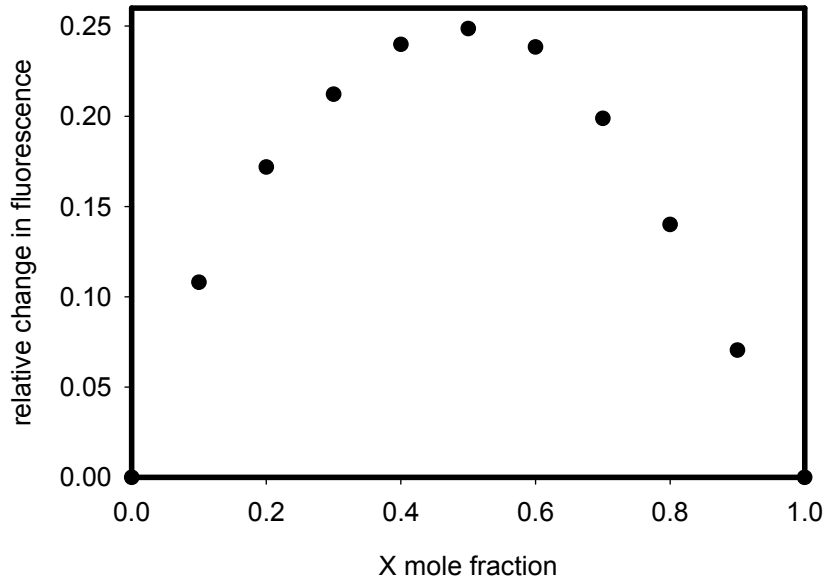
Supporting Information



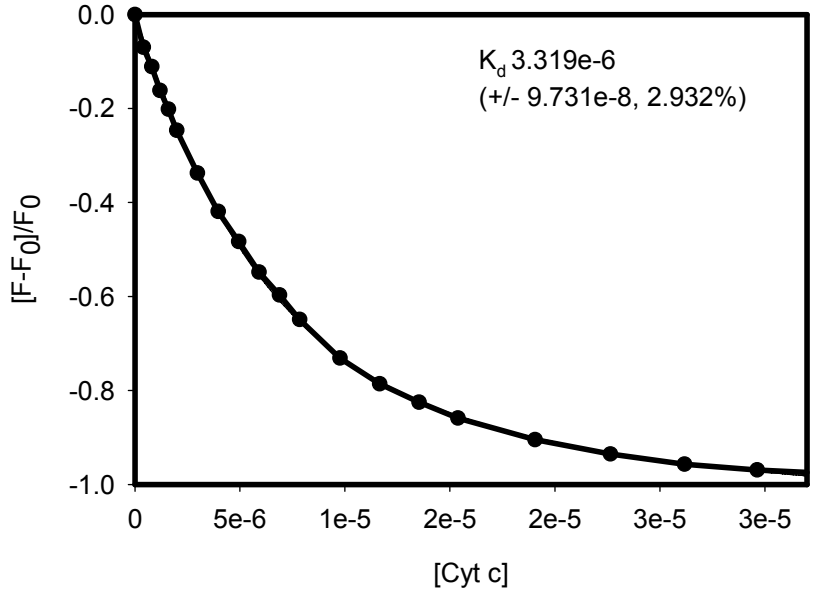
Job plot for binding of **1b** to Cytochrome *c*. Excitation wavelength was 330 nm; fluorescence emission was monitored at 401 nm as a function of the mole fraction of Cytochrome c. The total concentration of the two species was held constant at 20 μM (298 K, pH 7.4 and 5mM phosphate buffer).



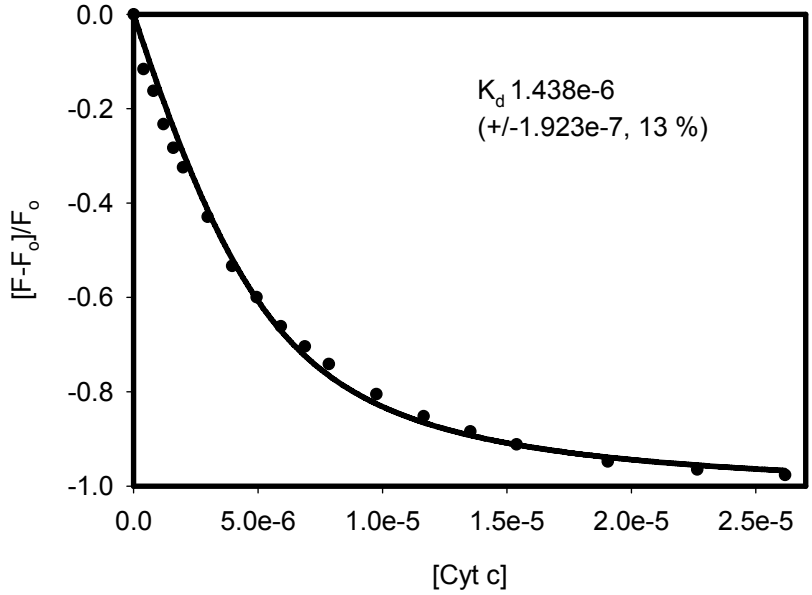
Job plot for binding of **1c** to Cytochrome *c*. Excitation wavelength was 330 nm; fluorescence emission was monitored at 401 nm as a function of the mole fraction of Cytochrome c. The total concentration of the two species was held constant at 10 μM (298 K, pH 7.4 and 5mM phosphate buffer).



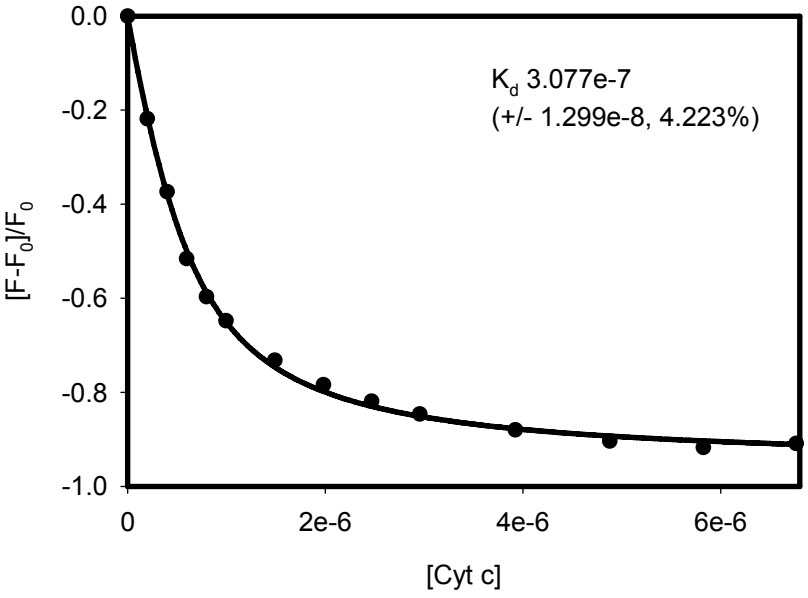
Job plot for binding of **1d** to Cytochrome *c*. Excitation wavelength was 330nm; fluorescence emission was monitored at 401 nm as a function of the mole fraction of Cytochrome c. The total concentration of the two species was held constant at 2 μM (298 K, pH 7.4 and 5mM phosphate buffer).



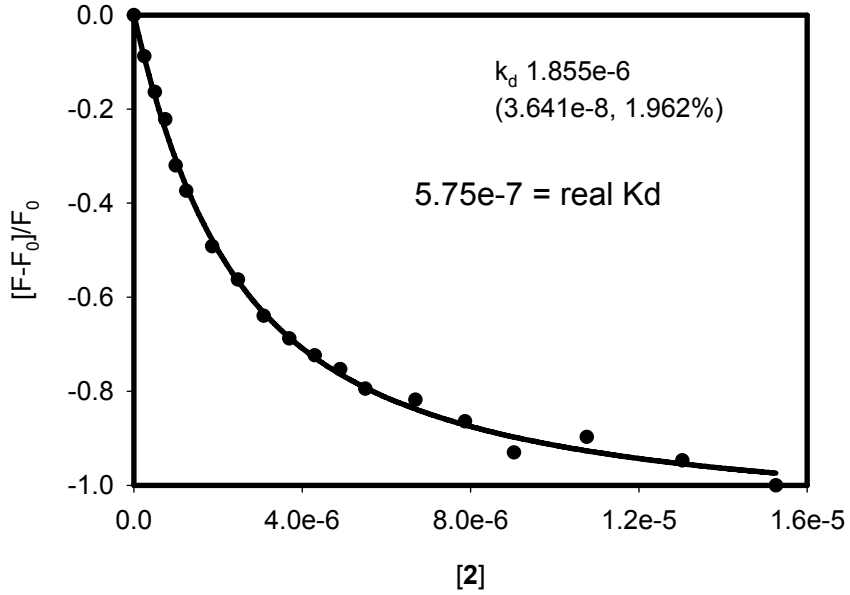
Fluorescence titration for the binding of 10 μM **1b** to Cytochrome *c*. (298 K, pH 7.4, 5mM sodium phosphate buffer). Excitation wavelength was 330 nm; fluorescence was monitored at 401 nm. Fit as a 1:1 complex.



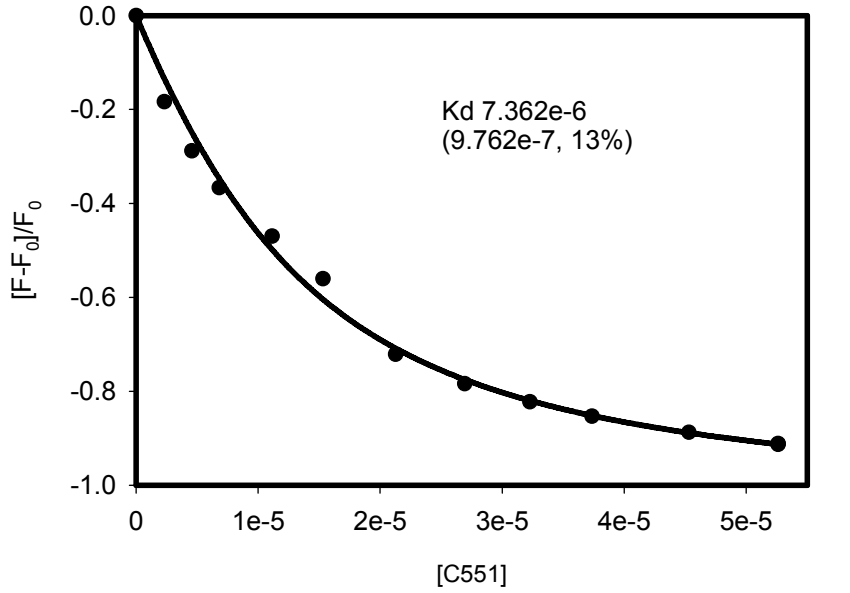
Fluorescence titration for the binding of 5 μM **1c** to Cytochrome *c*. (298 K, pH 7.4, 5mM sodium phosphate buffer). Excitation wavelength was 330 nm; fluorescence was monitored at 401 nm. Fit as a 1:1 complex.



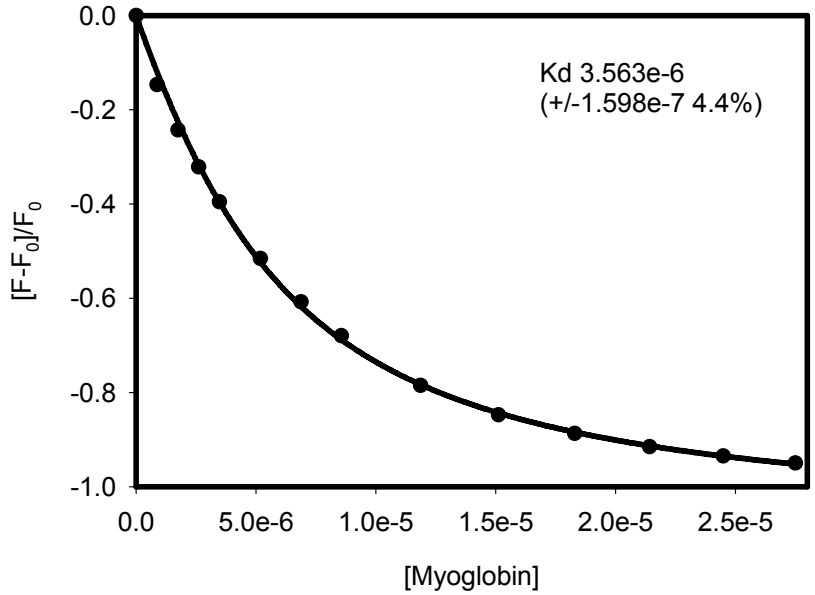
Fluorescence titration for the binding of 1 μM **1d** to Cytochrome *c*. (298 K, pH 7.4, 5mM sodium phosphate buffer). Excitation wavelength was 330 nm; fluorescence was monitored at 401 nm. Fit as a 1:1 complex.



Binding isotherm for the competition titration of **2** with 1μM **1a**:Cytochrome *c* (298 K, pH 7.4, 5mM sodium phosphate buffer). Excitation wavelength was 330 nm; decrease in fluorescence of **1a** was monitored at 401 nm until 1:1 stoichiometry with Cytochrome *c* and then the increase upon titration with **2** monitored to give the binding iostherm shown above.



Fluorescence titration for the binding of 4μM **1a** to Cytochrome c551. (298 K, pH 7.4, 5mM sodium phosphate buffer). Excitation wavelength was 330 nm; fluorescence was monitored at 401 nm. Fit as a 1:1 complex.



Binding isotherm for the titration of 4μM **1a** with myoglobin (298 K, pH 7.4, 5mM sodium phosphate buffer). Excitation wavelength was 330 nm; fluorescence was monitored at 401 nm. Fit as a 1:1 complex.