

Supplementary Materials

Fluorescence Detection of deoxyadenosine in *Cordyceps* spp. by Indicator Displacement Assay

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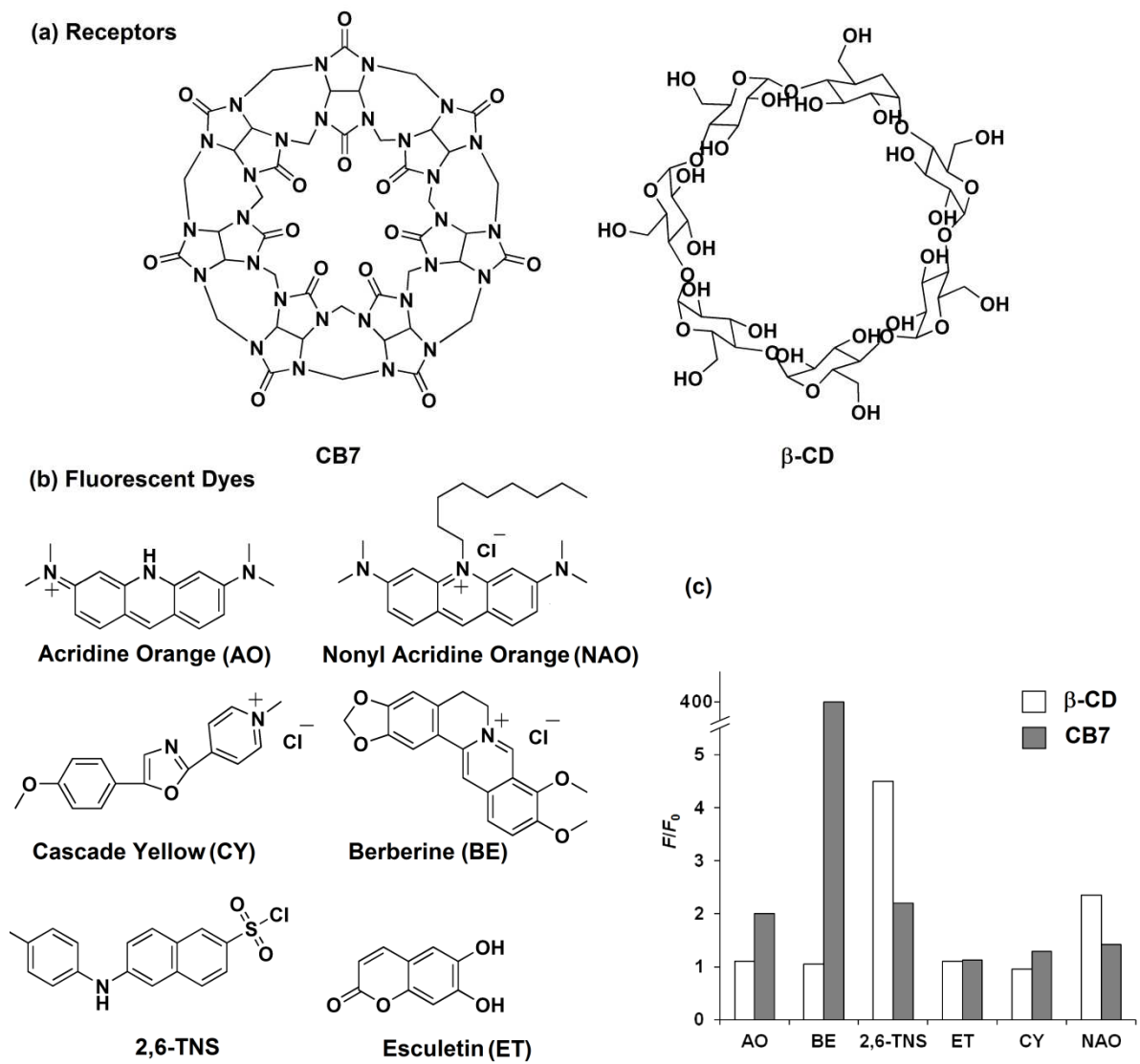
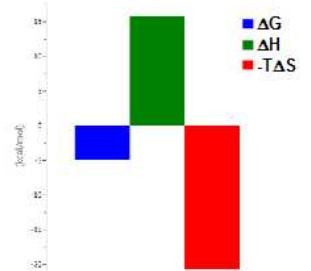
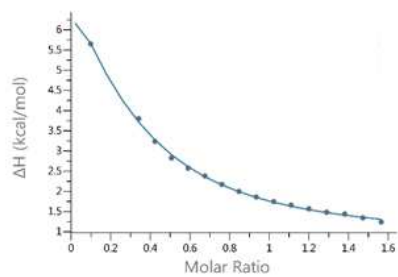
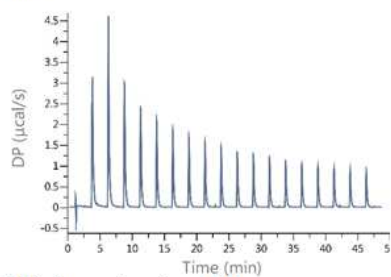
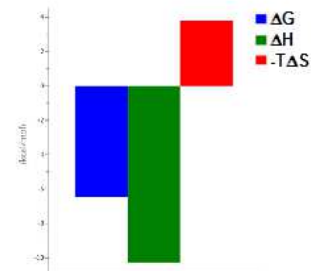
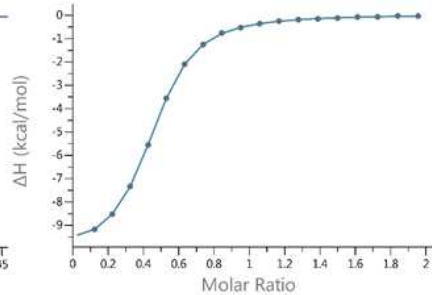
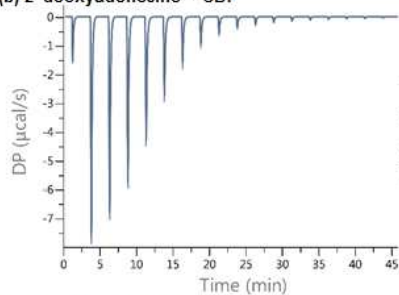


Figure S1. Chemical structures of macrocyclic receptors (a) and fluorescent dyes (b) tested in this study; (c) Fluorescent response of the dye without (F_0) and with 5 equivalent molar of β -CD (white) and CB7 (gray) in 10 mM ammonium acetate buffer (pH 5.0)

(a) AOH⁺ + CB7



(b) 2'-deoxyadenosine + CB7



(c) 3'-deoxyadenosine + CB7

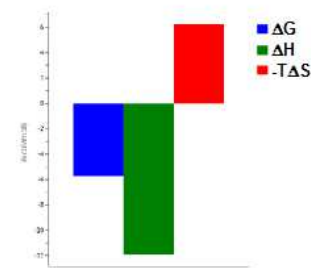
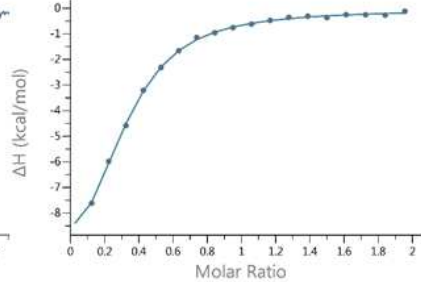
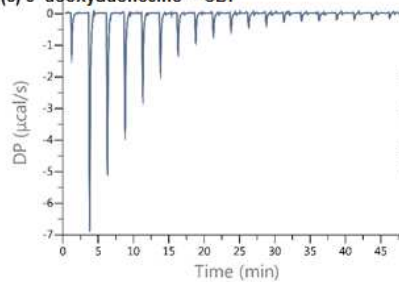


Figure S2. ITC analysis (MicroCal PEAQ-ITC, Malvern Instrument, UK) of CB7 (0.5 mM) and (a) AOH⁺ (4 mM), (b) 2'-deoxyadenosine (5 mM) and (c) 3'-deoxyadenosine (5 mM) in 10 mM ammonium acetate buffer (pH 4.0). Blue : ΔG , Green : ΔH , Red : $-\Delta S$ (kcal/mol).

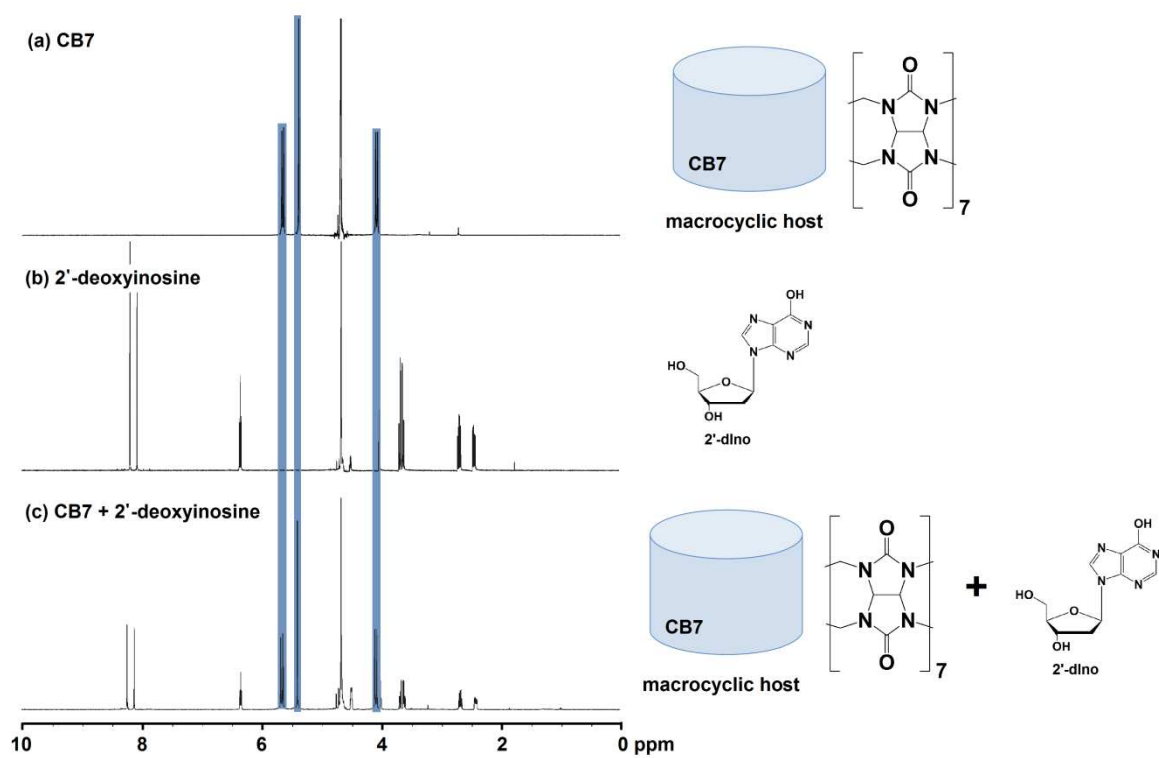


Figure S3. ¹H NMR (Bruker Ascend™ 400, D₂O) of the (a) CB7 (4 mM), (b) 2'-deoxyinosine (4 mM) and (c) the mixture between 2'-deoxyinosine and CB7 (4 mM each). Peaks of the CB7 were highlighted in blue.