

Supporting Information

Title: Highly efficient and selective extraction of uranium from aqueous solution by a magnetic device based on succinyl- β -cyclodextrin-APTES@maghemite nanoparticles

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Results

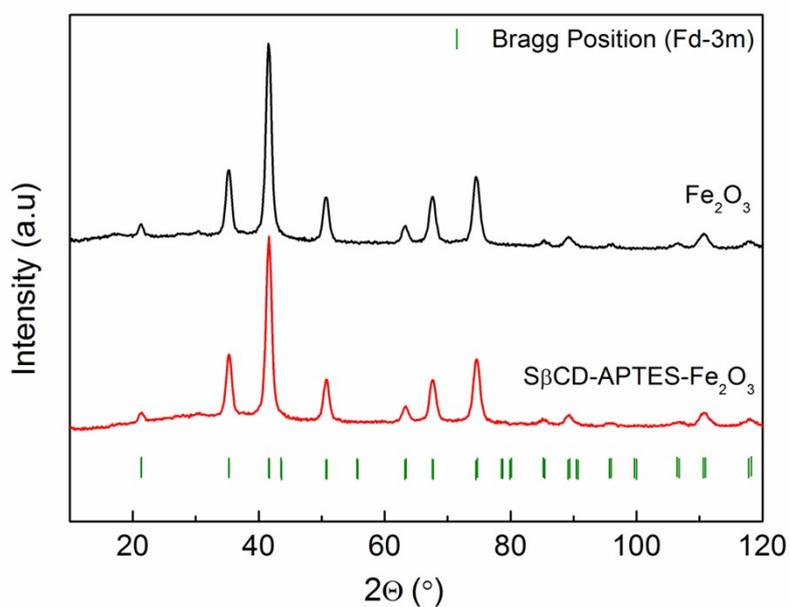


Figure S1: XRD pattern of Fe_2O_3 nanoparticles before and after surface functionalization with $\text{S}\beta\text{CD}$: (a) Fe_2O_3 ; (b) $\text{S}\beta\text{CD-APTES@Fe}_2\text{O}_3$.

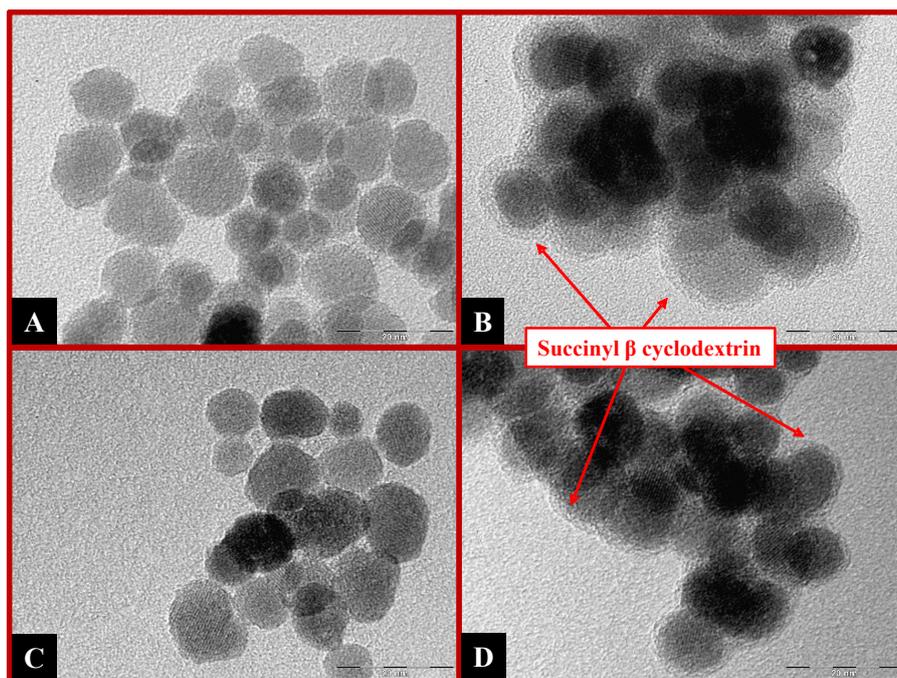


Figure S2: TEM images of APTES@Fe₂O₃ nanoparticles before and after surface functionalization with SβCD: (a), (c) APTES@Fe₂O₃; (b), (d) SβCD-APTES@Fe₂O₃.

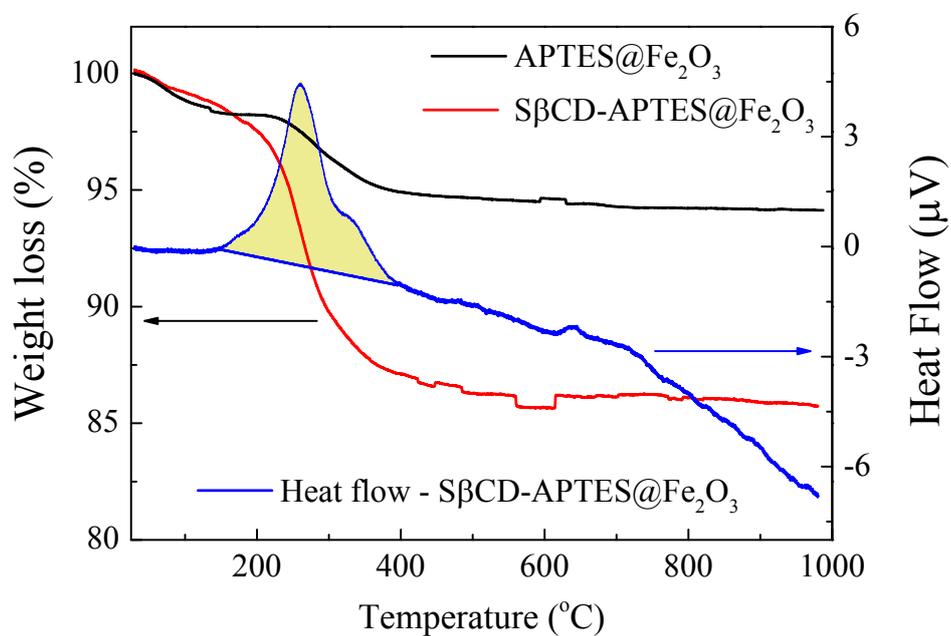


Figure S3: Thermogravimetric analysis of SβCD-APTES@Fe₂O₃ (red line) and APTES@Fe₂O₃ (black line).

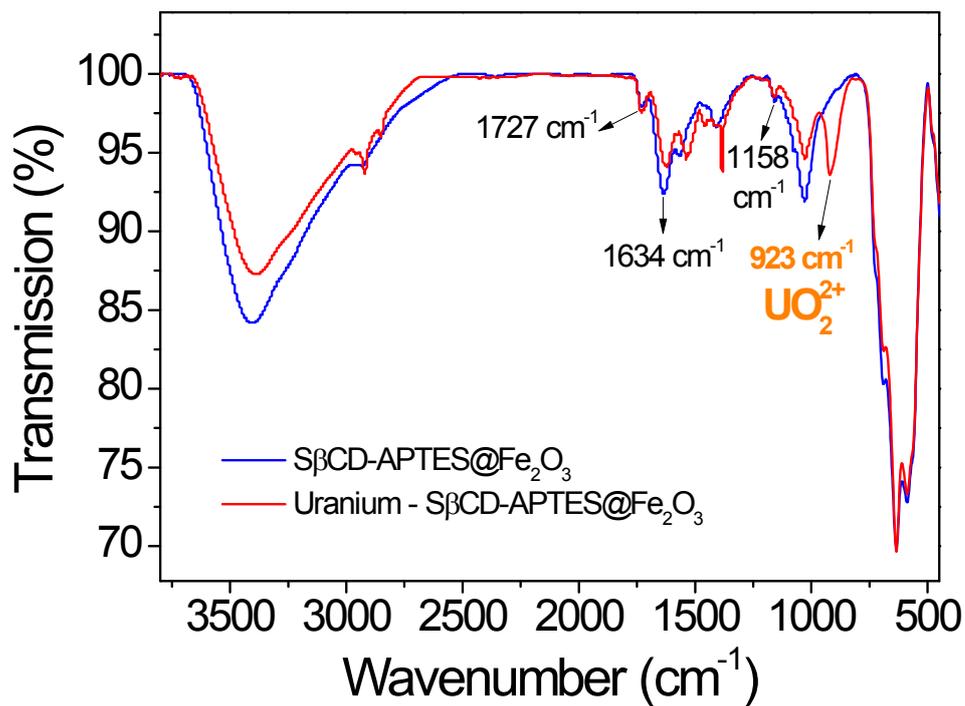


Figure S4: FTIR spectra of SβCD-APTES@Fe₂O₃: (blue) before U(VI) adsorption; (red) after U(VI) adsorption. Initial U(VI) concentration, 100 mg L⁻¹; adsorbent dose, 0.005 g; pH 6; 25 °C.

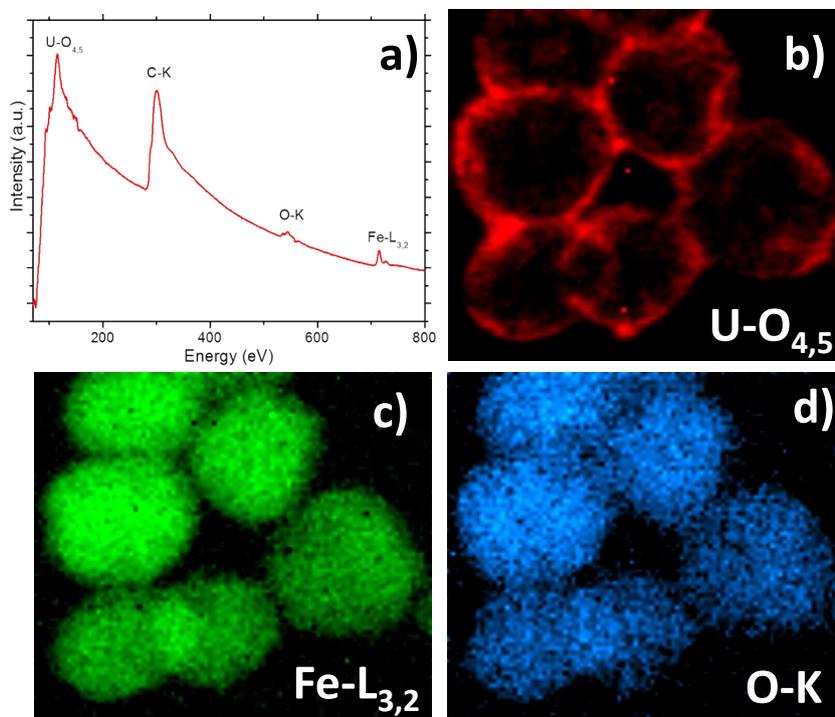


Figure S5: EELS extracted spectrum profile of U(VI)-loaded S β CD-APTES@Fe₂O₃ NPs. The spectrum image was recorded over the entire area with a resolution of 96 x 86 pixels, which results in a pixel size of 0.31 nm. The channel size for this experiment was set to 0.5 eV and the pixel time used was 0.1 s.

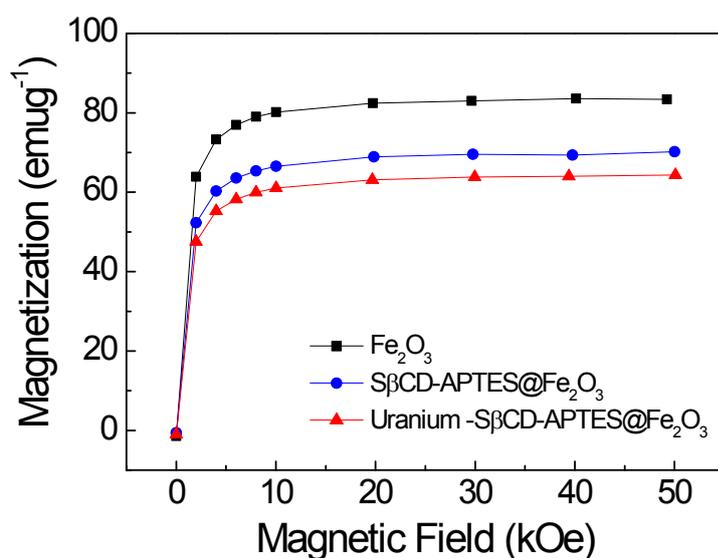


Figure S6: First emanation curves measured at 5 K, of raw NPs (black), and S β CD-functionalized NPs before (blue) and after complexation (red).

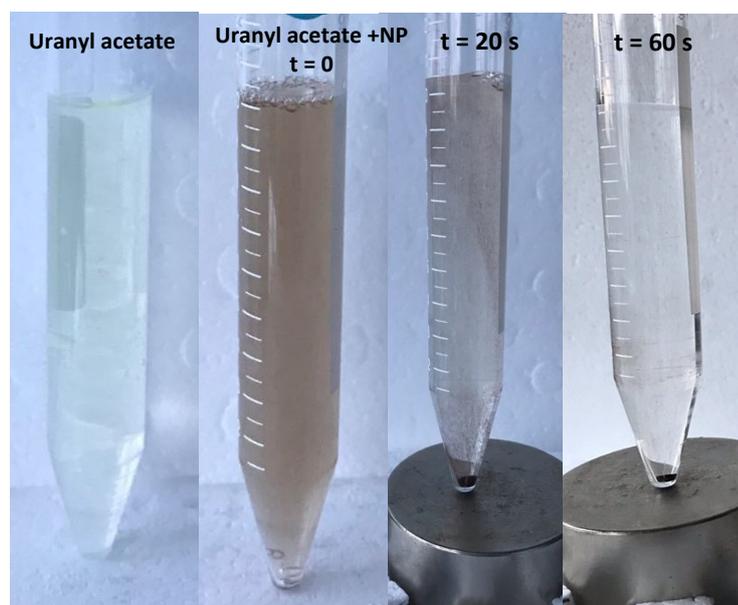


Figure S7: Example of magnetic sedimentation (t = 0, 20 and 60 s) of uranium-loaded S β CD-APTES@Fe₂O₃ nanoparticles by using a supermagnet (magnetic field = 1.3 T).