

Highly Porous and Monodisperse Magnetic Silica Beads

Prepared by a Green Templating Method

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Fig. S1: *Photograph of the MS beads series*

Fig. S2 : *DRIFT spectra of MA, MAS, and MS beads.*

Fig. S3 : *TEM images of microtome section of MAS beads*

Fig. S4 : *normalised UV-visible spectra of MeB in water and in HNO₃ aqueous solution, and of the supernatants after washing of the beads with HNO₃ aqueous solution.*



Fig. S1: Photograph of the MS beads series (each sample is characterized by a different value of $\Phi_{Fe_2O_3}$; from left to right $\Phi_{Fe_2O_3}$ increases from 0 to 2.1%)

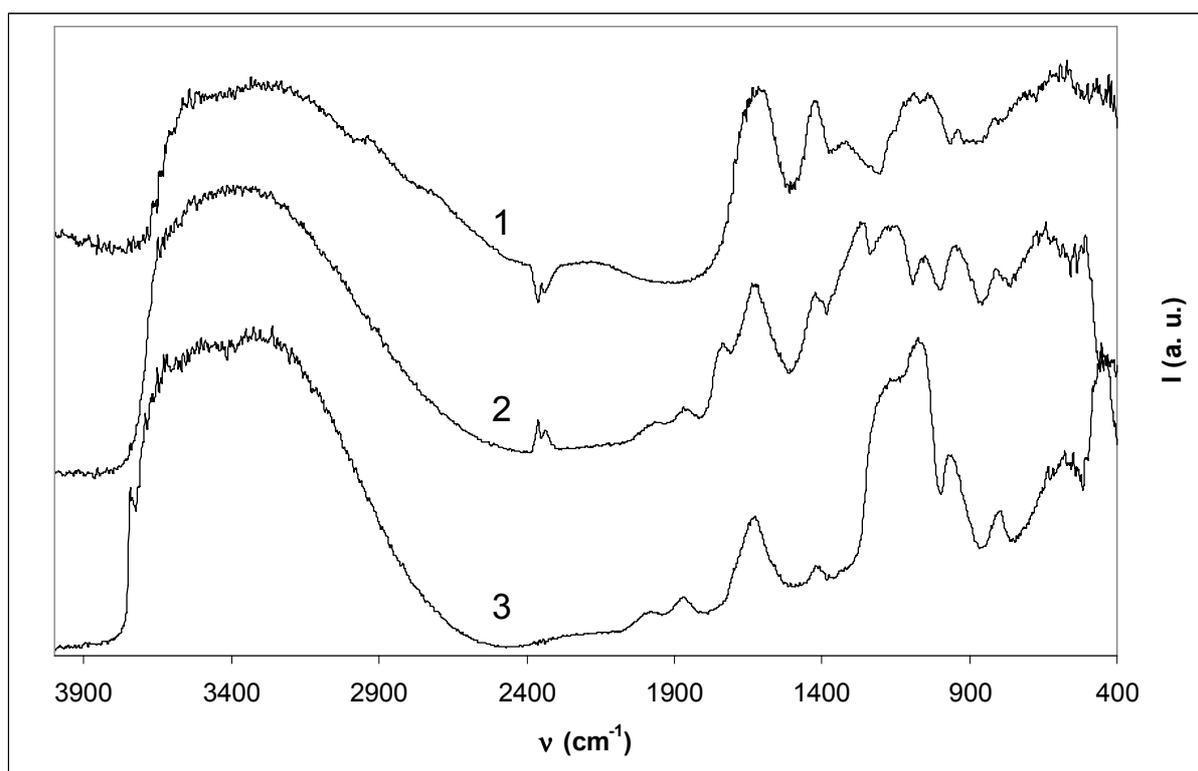


Fig. S2 : DRIFT spectra of MA (1), MAS (2) and MS (3) beads. ($\Phi_{\text{Fe}_2\text{O}_3} = 2.1\%$)

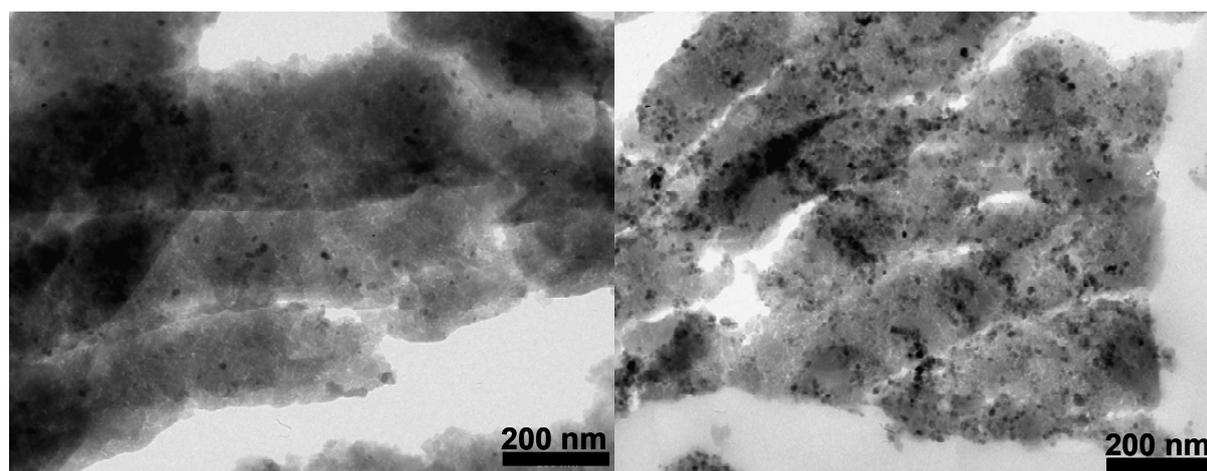


Fig. S3 : TEM images of microtome section of MAS beads (a) $\Phi_{Fe_2O_3}=0.06\%$;

(b) $\Phi_{Fe_2O_3}= 0.96\%$

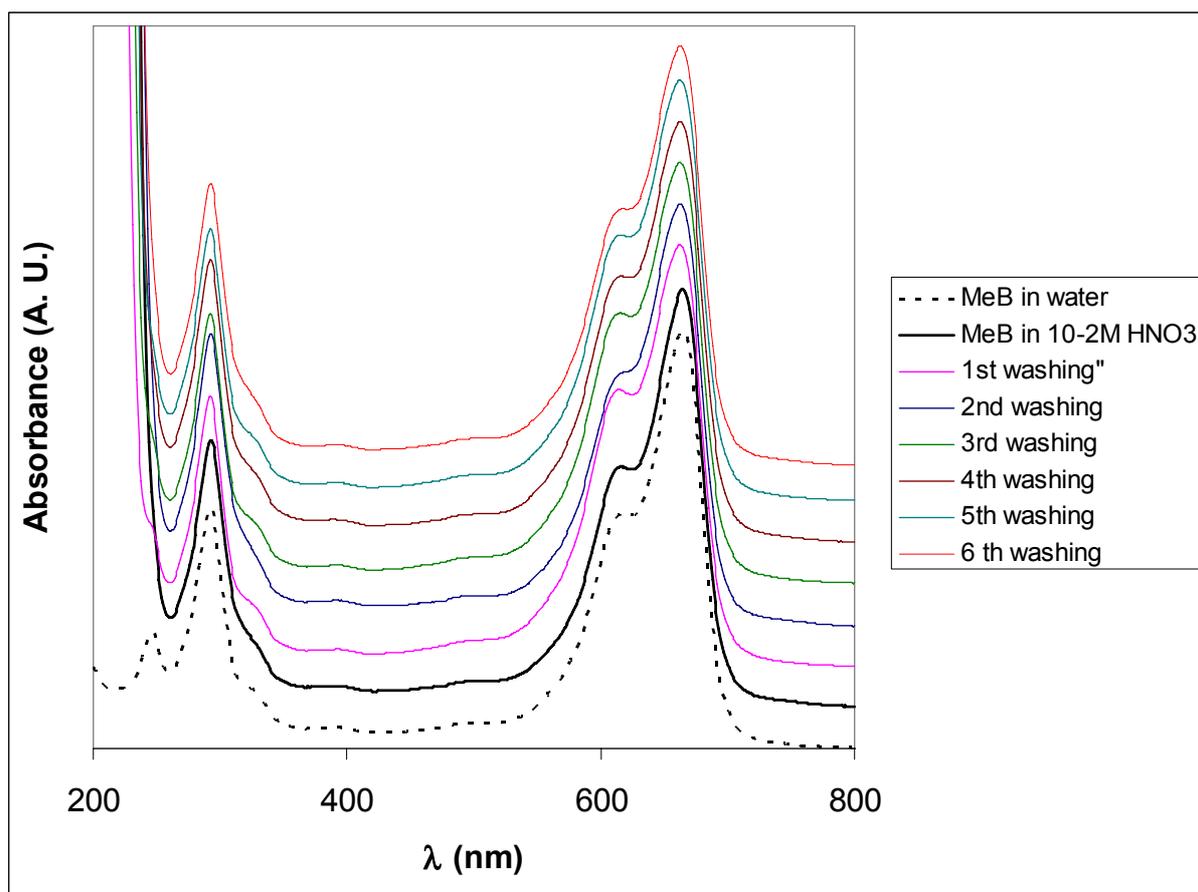


Fig. S4 : normalised UV-visible spectra of MeB in water and in 10^{-2} mol L $^{-1}$ HNO $_3$ aqueous solution, and of the supernatants after washing of the beads with 10^{-2} mol L $^{-1}$ HNO $_3$ aqueous solution.