SUPPORTING INFORMATION

Magnetic chelating nanoprobes for enrichment and selective recovery of metalloproteases from human saliva

Rui Oliveira-Silva, João Pinto da Costa, Rui Vitorino*, Ana L. Daniel-da-Silva*

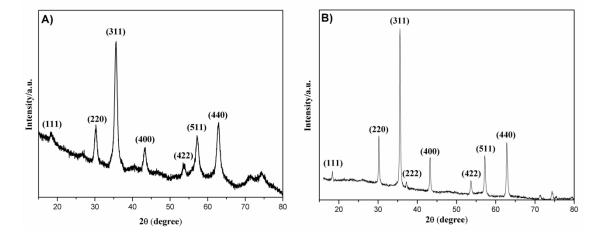


Figure S1 -A) X-ray diffraction pattern of 12 nm Fe₃O₄ nanoparticles. B) X-ray diffraction pattern of 50 nm Fe₃O₄ nanoparticles.

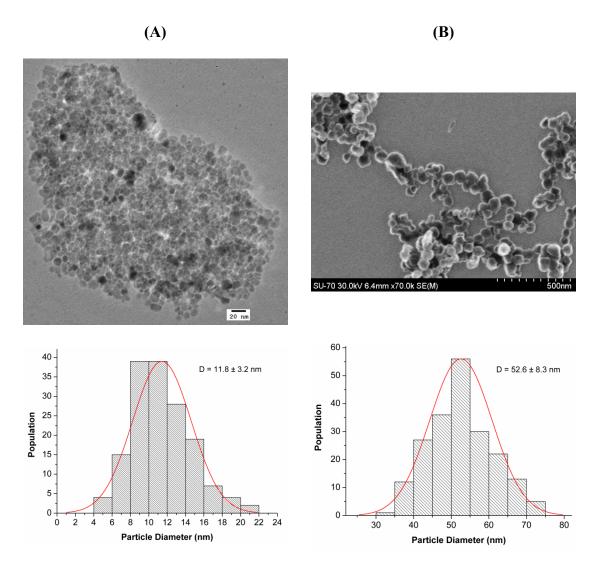


Figure S2 - Electron micrographs of (A) 12 nm Fe₃O₄ (TEM) and (B) 50 nm Fe₃O₄ nanoparticles (SEM) (top) and respective particle size histograms (bottom). The particle diameter was calculated as D = mean $\pm t$ s/ \sqrt{n} , where t is *t*-score (95% confidence), s is the standard deviation and n the sample size.

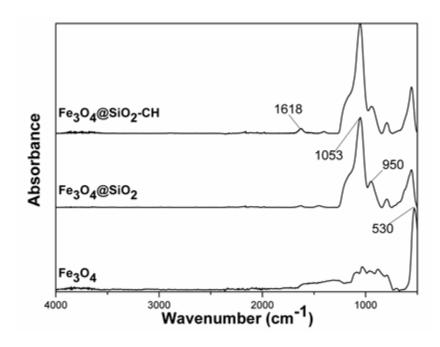
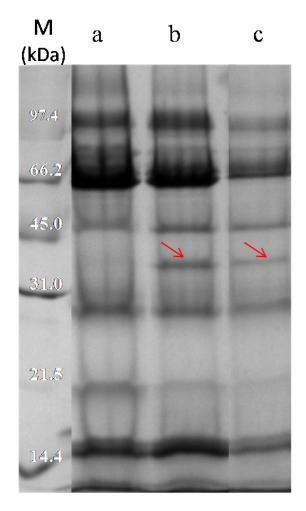


Figure S3 - ATR-FTIR spectra of 50 nm magnetic core NPs at distinct surface modification steps: Fe₃O₄, Fe₃O₄@SiO₂ and Fe₃O₄@SiO₂-CH functionalized at 70°C with 500 μ L of acetic acid during 24 hours.



 $\label{eq:Figure S4-SDS-PAGE (15\%) gel obtained after the incubation of a) 50Fe_3O_4@SiO_2-CH, b)12Fe_3O_4@SiO_2-CH and c) 12Fe_3O_4 NPs with human saliva. (M- Marker)$