Supporting Information

Effect of Bovine Serum Albumin on Tartrate Modified Manganese Ferrite Nano Hollow Spheres: Spectroscopic and Toxicity Study

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Figure S1. The comparative fluorescence intensity of BSA in presence of $T-MnFe_2O_4$ NHSs, H_2O and tartrate. Error bars are calculated from the standard deviation of 3 successive measurements for each case.

Continuous variation analysis (Job's plot)

Continuous variation analysis was done at 298.15 K (λ_{ex} = 329 nm). The fluorescence signal was recorded for solutions where the concentrations of both the BSA and the T-MnFe₂O₄ NHSs were varied while the sum of their concentrations was kept constant. The difference in fluorescence intensity (Δ F) of T-MnFe₂O₄ NHSs in the absence and presence of BSA was plotted as a function of the input mole fraction. Break point in the resulting plot corresponds to the mole fraction of the bound T-MnFe₂O₄ in the complex. The stoichiometry was obtained from [(1 – χ)/ χ], where, χ denotes the mole fraction of T-MnFe₂O₄ NHSs. The results presented are average of at least three experiments.



Figure S2. Job's plot depicting change in fluorescence intensity versus mole fraction of T-MnFe₂O₄ NHSs.

Table S2. The α -helical content (:	:3%) of BSA	for three successive me	easurements in different	concentration of T-MnFe ₂ O ₄ NHSs
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Conc. of T-MnFe ₂ O ₄	% of α-helix
NHSs (μg/ml)	
0	56.1
0.024	49.3
0.032	45.2
0.040	44.8
0.048	41.9



Figure S3. Analysis of the haematological parameters in T-MnFe₂O₄ NHSs treated rat. Changes in (a) W.B.C. count, (b) %W.B.C, and (c) Haemoglobin in rat blood treated with T-MnFe₂O₄ NHSs.



Figure S4. Antibacterial and antifungal activities of T-MnFe₂O₄ NHSs *in vitro*.