

Electronic Supplementary Information (ESI)

Hollow Spheres of Iron Oxide as an “Enzyme-mimic”: Preparation, Characterization and Application as Biosensors

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Figures:

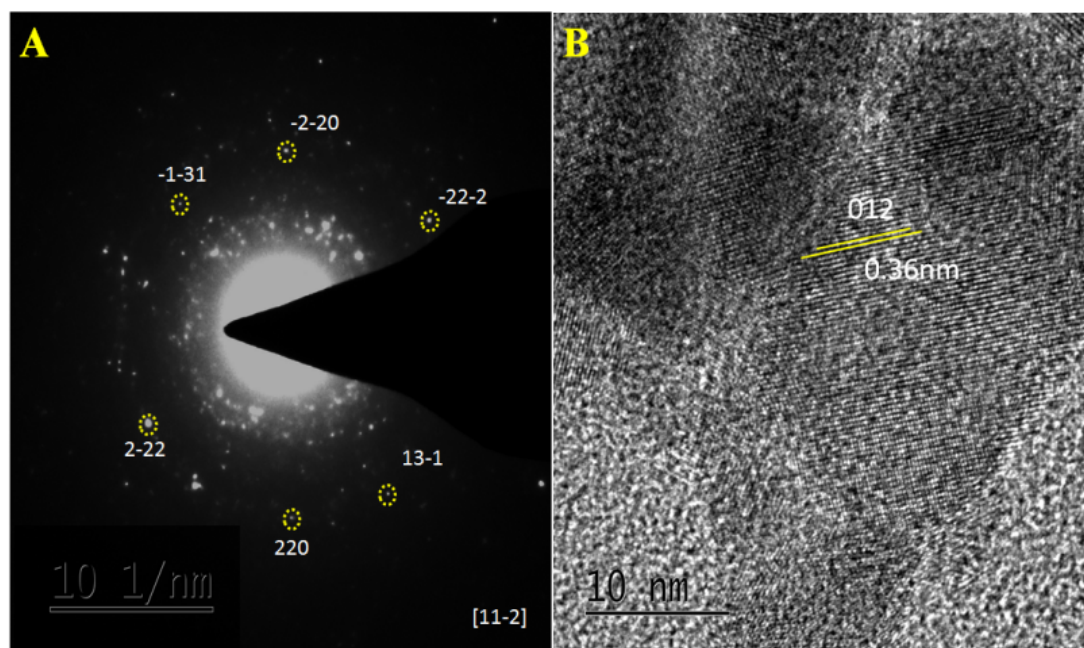


Figure S1: (A) Selected area electron diffraction and (B) diffraction fringes based TEM images of the synthesised HSFe_2O_3 nanoparticles.

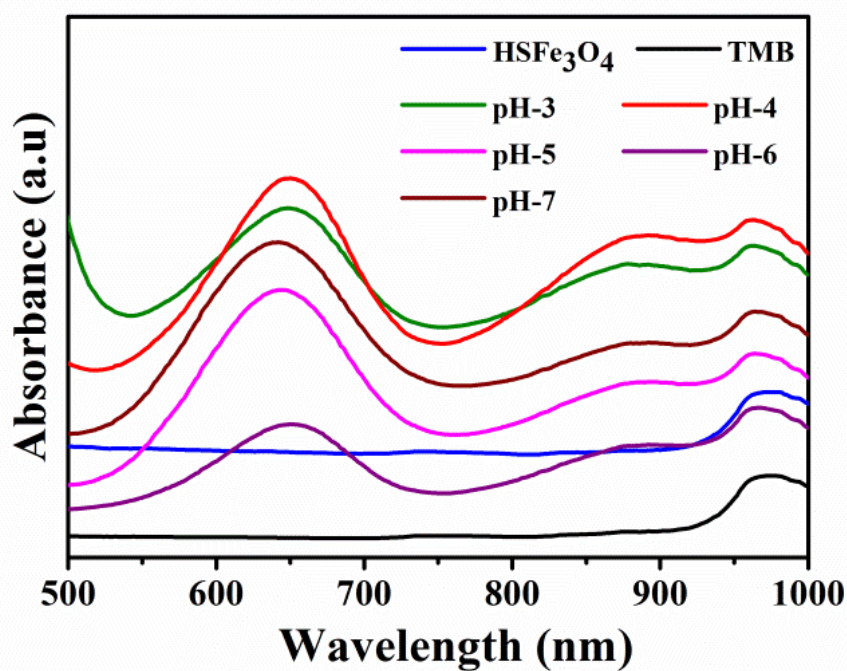


Figure S2: UV-visible spectra of TMB⁺ recorded at varied pH for peroxidase mimic assay.

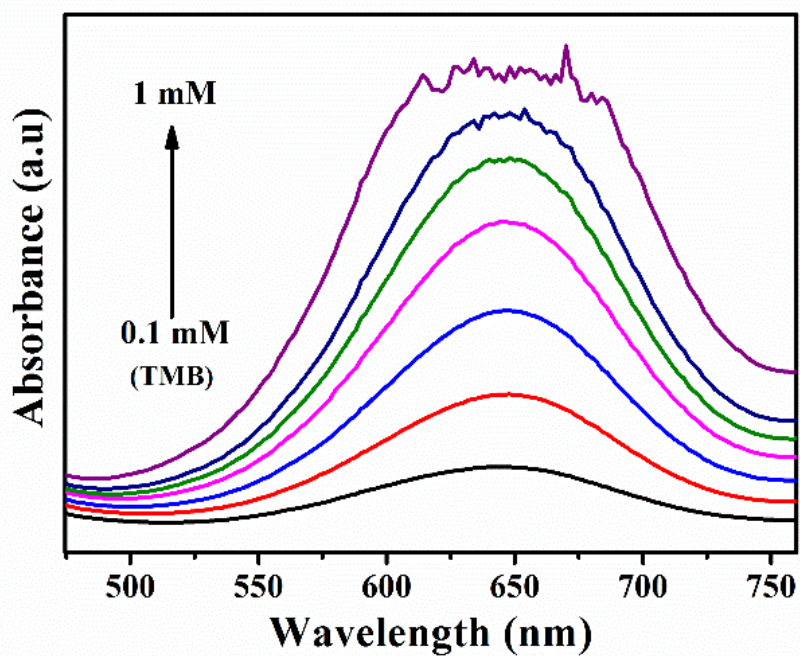


Figure S3: UV-visible spectra of peroxidase mimic assay recorded at varied TMB concentrations ranging from 0.1 mM to 1 mM. In this figure arrow indicates the direction of increasing concentration of TMB.

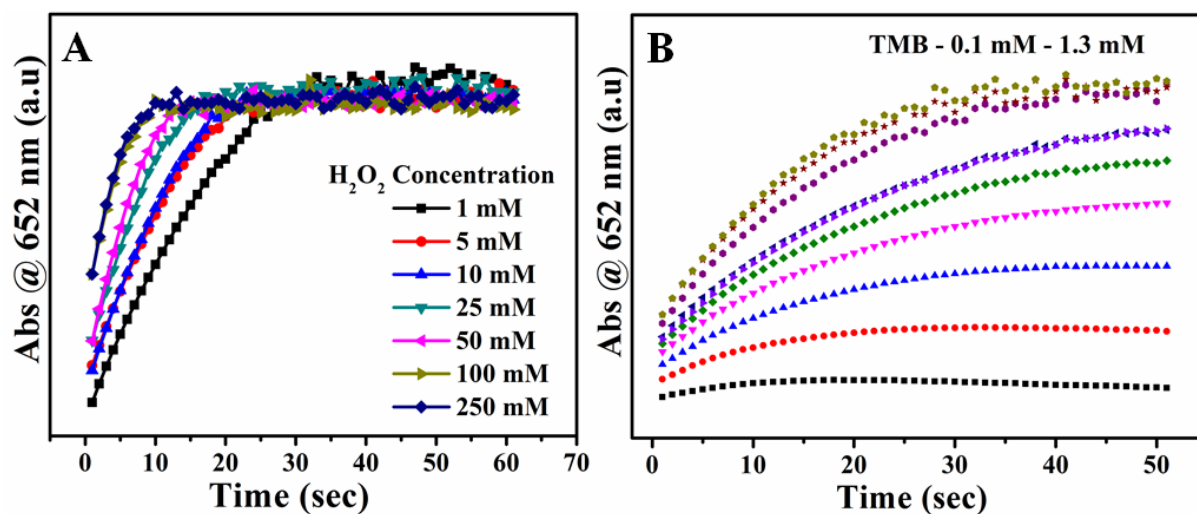


Figure S4: Time dependent UV-visible spectra of TMB – H₂O₂ system in presence of HSF₂O₃ (25 μg) and the corresponding absorbance values were measured at 652 nm. (A) TMB – 1 mM; H₂O₂ is varied from 1 mM to 250 mM and (B) H₂O₂ – 1 M; TMB is varied between 0.1 mM and 1.3 mM.

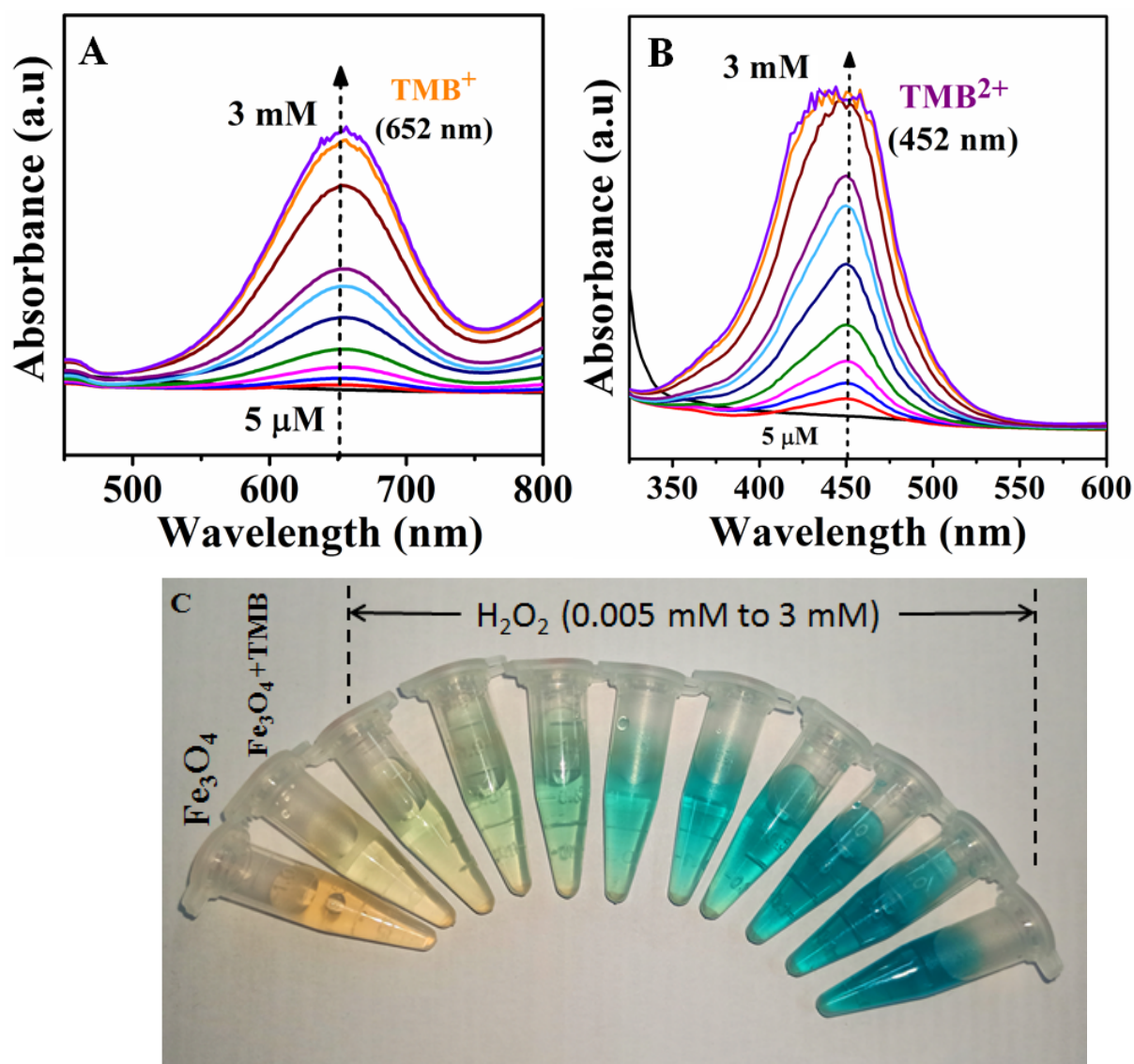


Figure S5: (A) and (B) UV-visible curves of TMB – H₂O₂ system in presence of HSF₂O₃ (25 μg) and TMB – 1 mM by varying H₂O₂ concentrations (from 5.0 μM to 3.0 mM). (A) 2 minutes reaction time; (B) After the reaction was arrested by adding H⁺ (20 μL of HCl, 0.01 M). (C) Photographs displaying the formation of blue color demonstrating the peroxidase mimic nanozyme using HSF₂O₃ in TMB – H₂O₂ system.

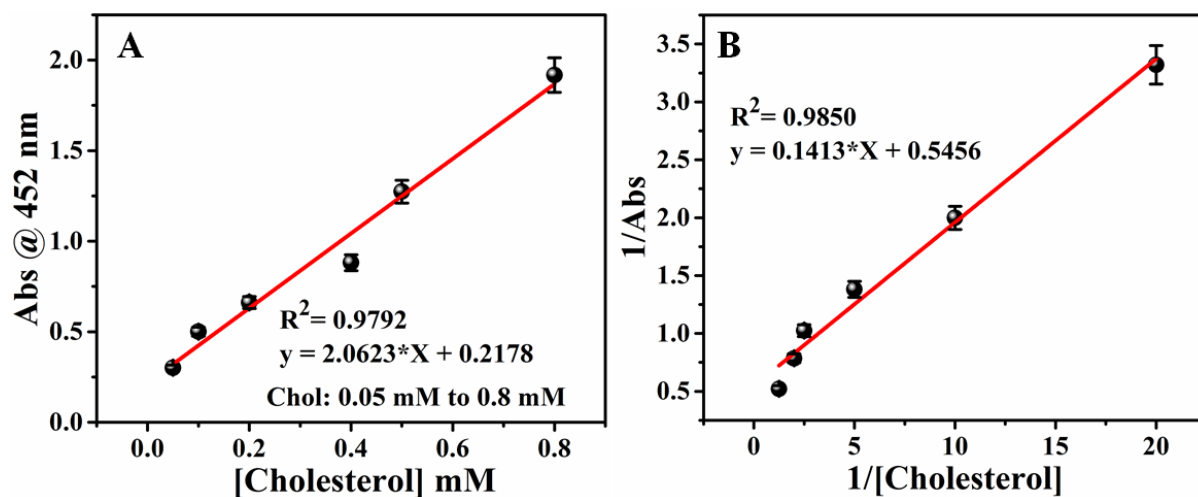


Figure S6: (A) UV-visible absorbance data recorded at 452 nm by varying the cholesterol concentration between 0.05 mM and 1.2 mM. (B) The calibration curves determined using Lineweaver–Burk equation. The error bars showed in this graph correspond to the standard deviation obtained from three independent measurements.