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A rationale for the rapid extraction of ultra-low-level uranyl ions in simulated bioassays regulated by Mn-dopants over magnetic nanoparticles†

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Although the sorption of uranyl ions and other heavy metal ions over magnetic nanoparticles is well reported, the parameters governing the sorption process over the magnetic nanoparticles have not been clearly enumerated. However, to increase the efficiency of the sorption over the surface of these magnetic nanoparticles, it is essential to understand the different structural parameters that are involved in the sorption process. The sorption of uranyl ions and other competitive ions in simulated urine samples at different pH was effectively accomplished over magnetic nanoparticles of Fe_3O_4 (MNPs) and Mn-doped Fe_3O_4 (Mn-MNPs). The MNPs and Mn-MNPs were synthesized using an easily modified co-precipitation method and were thoroughly characterised using several techniques, such as XRD, HRTEM, SEM, zeta potential, and XPS. The substitutional doping of Mn (1 to 5 at%) in the Fe_3O_4 lattice (Mn-MNPs) showed better sorption ability as compared to that of MNPs. The sorption properties of these nanoparticles were mainly correlated with the different structural parameters to understand the roles of surface charge and different morphological parameters. The interaction centres over the surface of MNPs with the uranyl ions were designated and the effects of ionic interactions with uranyl ions for these sites were calculated. Extensive XPS, *ab initio* calculations and zeta potential studies have provided deep insights into the different aspects that play key roles in the sorption process. These materials showed one of the best K_d values ($\sim 3 \times 10^6 \text{ cm}^3$) in a neutral medium with very low $t_{1/2}$ values (~ 0.9 min). The fast sorption kinetics (very low $t_{1/2}$) makes them amongst the best sorption materials for uranyl ions and optimal for the quantification of ultra-low-level uranyl ions in simulated bioassays.

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1. Introduction

The use of radionuclides has been greatly amplified due to the increasing demand for nuclear energy, application in

therapeutic and diagnostic purposes, and research and industrial application of radioisotopes. However, this can increase the probability of exposure of workers to radiation both externally and internally. External exposure can be controlled by (a) reducing the exposure time, (b) increasing distance from the source, and (c) applying adequate shielding. However, none of the above protective measures help to reduce the internal dose, *i.e.*, the Committed Effective Dose (CED). Hence, following internal contamination, the rapid determination of radionuclides is essential to decide on medical intervention, if required. Radionuclides emitting gamma rays or high-energy beta rays can be easily quantified by using *in vivo* monitoring techniques such as whole-body counting or organ-based counting (lung counting, thyroid counting, *etc.*). However, alpha-emitting and some beta-emitting radionuclides (*e.g.* ^{239}Pu , U_{nat} , ^{241}Am , ^{210}Po , ^{232}Th , and ^3H , ^{90}Sr) cannot be determined using the above-mentioned techniques. Depending on the chemical nature (solubility class) and biokinetic behaviour of these radionuclides, biological samples (urine, faeces, tissue, nasal swab, *etc.*) of the exposed person are analysed for the estimation of radionuclide intake and assessment of CED.

At present, the quantitative determination of alpha-emitting radionuclides from urine samples is labour-intensive and time-

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† Electronic supplementary information (ESI) available: Fig. S1: ED-XRF data for the different Mn-doped Fe_3O_4 system; (a) undoped Fe_3O_4 ; (b) Mn-1 (c) Mn-2 (d) Mn-3 (e) Mn-4 (f) Mn-5, Table S1: atomic weight percentage of elements as calculated from EDS data, Fig. S2: XPS spectra for Fe 2p for comparison of the process of sorption interaction; (a) Mn-2 before uranyl sorption and (b) Mn-2 after uranyl sorption post drying, Fig. S3: Plot for K_d versus the Mn-doping (at%) in the Fe_3O_4 lattice at neutral medium (pH-7). Table S2: SAED calculation for Fe_3O_4 system. Table S3: SAED calculation for Mn doped Fe_3O_4 system (Mn-3). Table S4: optimisation of the K-points for the bulk calculation of Fe_3O_4 . Table S5: comparison of cell parameters for Fe_3O_4 and Mn-doped Fe_3O_4 system. Fig. S4: optimised structures for (A) UO_2 and (B) UO_2 -ion. Table S6: energy and bond length for calculated UO_2 and $[\text{UO}_2]^{2+}$ ion system. (B) Calculation of interaction energy of $[\text{UO}_2]^{2+}$ ion with the Mn-doped Fe_3O_4 (3 1 1) surface. Fig. S5: speciation of uranyl ion at different pH (concentration of $\text{UO}_2^{2+} = 1 \text{ nM}$ and $\text{NO}_3^- = 3 \text{ M}$) as suggested in the Experimental section. The Figure is made by the Medusa Hydra-software plot. See DOI: <https://doi.org/10.1039/d3ra01957h>



consuming.¹ During normal situations, a large number of exposed individuals may need to be monitored for a short duration. Hence, novel approaches to detect, separate and estimate actinides from biological samples are being developed based on co-precipitation,² liquid-liquid solvent extraction,³ ion-exchange resins,^{4,5} extraction chromatography^{5,6} and nanotechnologies.^{7–10} The extraction of actinides using a solvent extraction technique is time-consuming and also generates organic waste. The most popular method is the separation of actinides by using ion exchange and extraction chromatography resins such as TEVA, UTEVA, TRU, *etc.* However, the separation process is comparatively slower and laborious. New strategies take advantage of nanoparticles, *i.e.* materials with nanoscale dimensions, which show high sensitivities to heavy metals and actinides, enhanced adsorption and responsiveness to external stimuli, such as magnetic fields.^{11–14} In the present study, the sorption of uranium over nanomagnetic materials like Fe₃O₄ and Mn-doped Fe₃O₄ was undertaken.

The removal of the different heavy ions from the environment along with the organic pollutants is a necessity for maintaining the ecological balance and protection of the environment and human health. The removal of the different heavy metals from wastewater has been effectively done by using different carbon nanoparticles^{15–18} and by photocatalysis.¹⁹ Recently, the removal of heavy metals using magnetic nanoparticles has also garnered substantial attention. Magnetic micro-nanoparticles have attracted considerable attention in several fields, such as analytical chemistry, wastewater treatment, catalysis, biochemistry medicine, *etc.*^{20–26} Sorbents comprised of iron oxides like magnetite (Fe₃O₄) and maghemite (γ -Fe₂O₃),^{27–30} with different examples for other iron oxide phases, may be attached to different ligands or polymeric materials,^{31–35} and have been utilised for extracting the toxins mainly from wastewater. These materials are good contenders as the oxides are mostly non-toxic materials and they are easily separated by magnetic separation, which is fast and effective. One of the most important properties of magnetic nanoparticles (MNPs), which makes them suitable for use as compared to other materials, is their magnetic behaviour. Bare MNPs such as Fe₃O₄ are dispersible in water because of the repulsion between particles due to like-charges on the surface, their small size and they exhibit no magnetic properties in the absence of a magnetic field. However, under an external magnetic field, these MNPs tend to agglomerate because of magnetic dipole interactions. Since these nanoparticles have a high surface area, as well as good pore size, many metal ions and dyes can be sorbed on the surface and occupy pores or lattice sites of nanoparticles.^{36–40} A plethora of nanoparticles, magnetic and using different supports, have been utilised for the sorption of actinide materials like uranyl ions from the solution.^{41–43} Such incorporated/adsorbed NPs can be attracted towards the magnetic field and can be used for the removal of toxic materials from the environment as has been discussed for the separation of ultra-low level (ULL) uranyl ions present in urine following internal contamination. The idea is not only to adsorb the ULL-uranyl ions but also to separate them

magnetically, with the rapid separation process making it tenable for the estimation procedure.

Mn-doped Fe₃O₄ has garnered a lot of attention lately due to the different credible properties of these doped magnetic materials. Precise control over the Mn dopant included in the iron oxide lattice should permit the tailoring of the sorbent reactivity towards the analytes of interest. These materials have been utilised lately for several purposes like the sorption of several cationic dyes,⁴⁴ sorption of heavy metals like Hg,⁴⁵ and CO adsorption with different dopant segregation.⁴⁶ Similarly, functionalised Fe₃O₄ and Mn-doped Fe₃O₄ have been used for simultaneous dye and metal ion ultrasound-assisted adsorption.⁴⁷ The Mn-Fe₃O₄ system dispersed over reduced graphene oxide has been utilised as a photo-Fenton agent under neutral pH to degrade certain dyes like Rhodamine-B with high efficiency.⁴⁸ Warner *et al.* have shown a suitable method for the synthesis of Mn-doped Fe₃O₄ using commercial magnetite nanopowder where they have shown that the level of Mn dopant in the ferrite has a significant effect on the reactivity of the materials toward selected heavy metal analytes (*e.g.*, Co, Ni, Cu, Zn, As, Ag, Cd, Hg, and Tl).⁴⁹ Though there have been several reports, the exact parameters governing the sorption for the doped Fe₃O₄ systems have not been completely revealed. If the roles of the different structural parameters along with that of the solvent are known properly, it will be easier to tune the sorption properties for any MNPs.

Following accidental internal contamination (with uranium), the rapid determination of radionuclides is essential to decide the requisite medical intervention. Therefore, to separate the uranyl ions from the urine bioassays using magnetic separation, a very efficient sorbing material with very fast kinetics is needed. To achieve this goal, the sorption of uranium on nanomagnetic materials like Fe₃O₄ and Mn-doped Fe₃O₄ was initially undertaken. The effects of the pH on the doped Mn-Fe₃O₄ surface, as well as the effect of Mn-doping over Fe₃O₄ for the sorption of the uranyl ions, were studied. Also, the different structural parameters that directly regulate the process of sorption over these Mn-MNPs and MNPs along with the particular sites where they were sorbed were investigated using XPS, and a theoretical understanding was formulated to substantiate the sorption efficiencies of these materials. The MNPs and Mn-MNPs having very strong sorption properties (high K_d values) and simultaneously very fast kinetics for the process of sorption (low $t_{1/2}$ values) can be very conveniently used for the separation of low-level uranyl ions from the different bioassays.

Conventional methods that are presently used for the bioassay sample analysis of ultra-low-level (ppb) actinide (U, Pu, Am *etc.*) detection involve acid digestion, co-precipitation, ion-exchange chromatographic separation, alpha source preparation and activity estimation using an alpha spectrometer. These processes take nearly 4–5 working days for single sample analysis. Thus, to develop a faster bioassay method, it is important to separate actinides with faster kinetics and with selective sorption over the substrate. The above MNPs and Mn-MNPs will effectively lead to a paradigm shift for the development of



a faster estimation process, which includes selective sorption as the major focus.

2. Experimental methodologies

2.1 Synthesis of nano Fe_3O_4 (MNPs)

Nano Fe_3O_4 was synthesized *via* a modified co-precipitation route^{50,51} using $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ as a precursor for Fe^{3+} and $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ as a precursor for Fe^{2+} in stoichiometric ratios. Here, 100 mL of 0.1 M $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$ solution was taken in a 3-necked round bottom flask, and then 50 mL of 0.1 M $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ were added. N_2 gas was used to purge the solution, with stirring for 1 h, to remove the dissolved oxygen. Next, 100 mL NH_4OH solution was added to the solution dropwise from a burette under an inert atmosphere. The mixture was stirred for 3 h at room temperature. The resulting black particles were washed several times with distilled water followed by acetone. The products were separated magnetically and dried at $\sim 60^\circ\text{C}$ for 5–6 h.

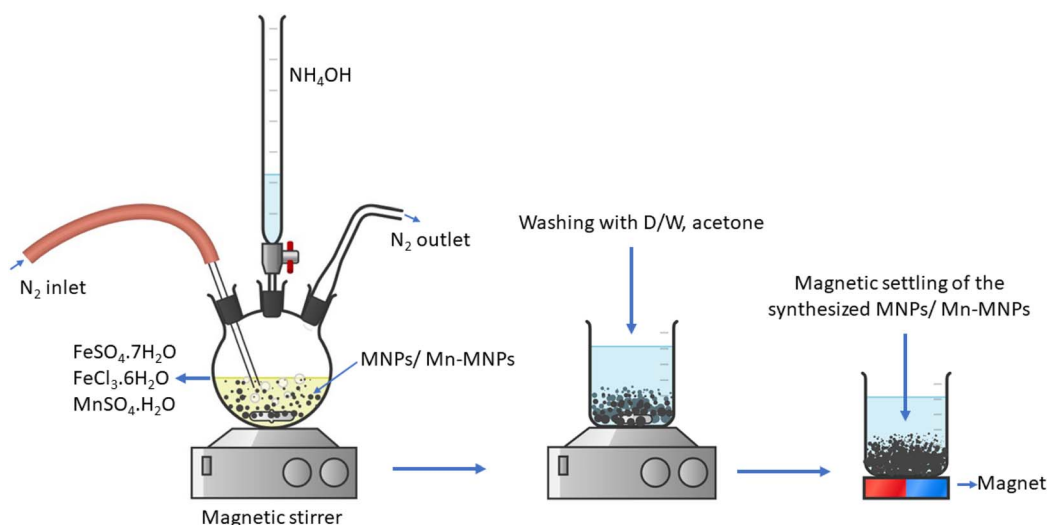
2.2 Synthesis of Mn-doped Fe_3O_4 (Mn-MNPs)

The as-synthesized nano Fe_3O_4 was then doped with Mn to improve its sorption property. The Mn was doped in atom percentages, where a stoichiometric amount of $\text{FeSO}_4 \cdot 7\text{H}_2\text{O}$ was substituted by $\text{MnSO}_4 \cdot \text{H}_2\text{O}$ (0.1 M) solution added to 0.1 M $\text{FeCl}_3 \cdot 6\text{H}_2\text{O}$. Different stoichiometric amounts of $\text{MnSO}_4 \cdot \text{H}_2\text{O}$ (0.1 M) were added so that the Mn-doped percentages were 1, 2, 3, 4 and 5 at% in the Fe_3O_4 lattice. The required amount of N_2 gas was used to purge (Scheme 1) the solution, with stirring for 1 h, to remove the dissolved oxygen. Next, 200 mL NH_4OH solution was added to the solution dropwise from a burette under an inert atmosphere. The mixture was stirred for 2.5 h. A black precipitate was obtained, which was washed several times with distilled water followed by acetone, as shown in Scheme 1. The black powder was separated magnetically and dried. The Mn-doping percentage ranged from 1 to 5 at% and these

samples were labelled as Mn-1, Mn-2, Mn-3 Mn-4 and Mn-5, respectively, and are collectively denoted as Mn-MNPs.

2.3 Characterization

Crystallographic phase analysis was carried out from powder X-ray diffraction (XRD) measurements using a Phillips Analytical Diffractometer with Ni-filtered $\text{Cu K}\alpha$ radiation. The diffractograms were recorded in the $10\text{--}80^\circ$ (2θ) region. The BET surface area, pore volume, and pore size distribution profiles of different samples were obtained from the physical adsorption of N_2 at 77 K, using a Micromeritics ASAP 2020 analyser. About 100 mg of a sample was utilized for this purpose, which was degassed under vacuum (10^{-6} Torr) at 300°C , before N_2 adsorption. Raman spectra were obtained using a 632.8 nm line of a diode laser for excitation and the scattered light was analyzed using a 0.9 nm single-stage monochromator coupled with a CCD detector. Laser Raman spectra were taken on a HORIBA Jobin Yvon instrument. The particle size, structure parameters and morphology were investigated using a TEM (JEOL F2100) equipped with a LaB_6 emitter, Thermo Noran Energy Dispersive X-S4 ray Spectroscopy (EDXS) system and a Gatan Model 666 Parallel Electron Energy Loss Spectrometer (PEELS) system. X-ray Photoelectron Spectroscopy (XPS) study was performed using a Thermo Fisher Scientific NEXSA instrument operating at 12 kV anode voltage with 6.50 mA filament current (1486.6 eV Al $\text{K}\alpha$ dual anode source). The data was taken at a pass energy of 50 eV under vacuum at 9×10^{-8} mbar. As an internal reference for the absolute binding energy, the C 1s peak (284.5 eV) was used. All the deconvolutions are made by the CASA software with a Voigt-type peak having GL 75 to GL 30 without imparting any asymmetry. The baseline was made using the Shirley function. Zeta potential measurements were made at 25°C using a quartz cuvette with a Malvern Zetasizer nanoseries, by phase analysis light scattering with an applied field strength of $2.5 \times 10^3 \text{ V m}^{-1}$. The light source was a He–Ne laser (632.8 nm) operating at 4.0 mW. About 0.5 mg



Scheme 1 The synthesis of nano Fe_3O_4 and Mn-doped Fe_3O_4 samples.

mL⁻¹ of the sample was dispersed in 1 mL of water. The surface morphologies and particle size were studied using a Seron AIS 2100 Scanning electron microscope. The dc magnetization measurements were carried out using a Vibrating Sample Magnetometer (VSM, Cryogenic Ltd, UK), as a function of both temperature and magnetic field. Magnetization as a function of the magnetic field was recorded at ~298 K (RT) over an applied magnetic field of +5 to -5 kOe for all samples. The elemental detection for the Mn-doped Fe₃O₄ samples along with the different competing ions of Na⁺, K⁺, Ca²⁺ and Mg²⁺ were done with the ICP-OES technique with a Thermo Fisher ICP-OES-iCAP-6500 instrument with a CID detector. A light-emitting diode (LED) fluorimeter (model: UA2, make: Quantalase Enterprises Pvt Ltd Indore, India) was used for the fluorescence measurement of uranium in an aqueous medium at the excitation and emission wavelengths of 405 and 516 nm, respectively.

2.4 Computational details

Quantum chemical calculations were performed using density functional theory. Here, the Vienna *ab initio* simulation package (VASP)⁵² electronic structure code was used along with the projector augmented wave (PAW)⁵³ package. The doped systems were constructed using 56 atoms with the Fe₃O₄ crystal structure as shown in Fig. 15. Geometry optimisation was carried out by relaxing both cell parameters and ionic positions under the framework of the generalized gradient approximation (GGA) using the Perdew–Burke–Ernzerhof (PBE) functional.⁵⁴ Brillouin zone sampling was carried out using the Monkhorst and Pack scheme⁵⁵ with a *k*-point mesh size of 3 × 3 × 3. Self-consistent iteration was continued until energy convergence reached 10⁻⁷ eV. The *E*_{cut-off} value was kept fixed at 500 eV throughout the calculation. The bulk calculation for the Fe₃O₄ was performed using the ICSD-029129, which matches the Fe₃O₄ bulk structure as shown in Fig. 1a. The surface calculation was made by cutting the (3 1 1) surface from bulk Fe₃O₄ with 112 atoms with O atoms facing the surface. The surface calculation for the Mn-doped Fe₃O₄ was made from the octahedral Mn-doped Fe₃O₄ (3 1 1) with 112 atoms. The surface calculations were done with a *k*-point mesh size of 1 × 1 × 1. The calculations were done with a vacuum space of 20 Å over the surface and with a maximum cut-off potential of 500 eV. The convergence threshold was set to 10⁻⁵ eV for electronic optimisation and the force convergence was set to 0.02 eV Å⁻¹ for structural optimisation. The interaction with the uranyl ion was done using a total of two electrons, less than the total valence electrons of uranyl and uranyl adsorbed over Mn, Fe (Oct), and Fe(tet) in the above surface. The interaction energy of the surface and the uranyl ions was calculated with K points K (1 1 1) along with the same vacuum region above the surface.

2.5 Sorption studies over magnetic nanoparticles

A known amount of U(nat.) standard (~75 ppb) was spiked in a conical flask and then dried on a hot plate. Next, 2.5 mL conc. HNO₃ was added and evaporated to dryness, then 5 mL of 0.1 M HNO₃ was added and after 10 minutes, 45 mL of distilled water

(D/W) was added to prepare 50 mL of known U(nat.) standard reference solution A. 2 mL fraction from solution A was analyzed using the LED fluorimeter for the initial concentration of U(nat.) present in the standard solution A. Next, ~50 mg of magnetic nanosorbent (MNPs/Mn-MNPs) was added to the standard solution A with sonication. At different time intervals, 2 mL fractions of the supernatant solution were collected to determine the residual uranyl ion concentration. Before collection of the supernatant fraction, all samples were magnetically settled. These fractions were analyzed in a LED fluorimeter (1 mL sample solution + 5 mL sodium pyrophosphate buffer solution) to determine the U(nat.) concentration present in the respective aliquots after adsorption on the magnetic nanoparticles. An equivalent study was carried out at different pH (3, 7 & 10) for all the MNPs and Mn-MNPs samples. pH adjustment of the sample solutions was done using ammonium hydroxide solution.

A 1.5 L stock aqueous solution, containing Na (1420 µg mL⁻¹), K (814.7 µg mL⁻¹), Ca (44.2 µg mL⁻¹), and Mg (71 µg mL⁻¹) ions, was prepared, mimicking the average concentration range of these ions present in the bioassay sample urine.⁵⁶ A natural uranium standard (prepared in 3 M HNO₃ medium; the concentration of uranyl ions in the standard solution was ~9.4 µg mL⁻¹) was spiked (1.5 mL) in the 100 mL stock solution.

To the 100 mL stock solution, ~100 mg of magnetic nanosorbent was added with sonication. The concentrations of the different ions (Na⁺, K⁺, Mg²⁺ and Ca²⁺) present in the residual solution at different time intervals were studied with ICP-OES. The effects of competitive ions on the sorption over Fe₃O₄, Mn-2 and Mn-4 were studied at acidic and basic pH (pH = 4 & pH = 9, respectively).

To understand the kinetics of uranyl ion sorption over the different MNPs and Mn-MNPs, the data were mainly fitted with a sigmoidal curve in Origin 8.0. The fitting with the sigmoidal curve signified a 1st order reaction. The adj. *R*-square (signifying the statistics for the fitting of the curve) varied from 0.96047 to 0.99826, which signified a good fit.

3. Results and discussion

3.1 X-ray diffraction (XRD)

Fig. 1A shows the characteristic powder-XRD patterns for samples studied in the present investigation: bulk Fe₃O₄, nano Fe₃O₄ and the Mn-doped Fe₃O₄ (Mn-1, Mn-2, Mn-3, Mn-4 and Mn-5). The XRD pattern corresponding to nano-Fe₃O₄ (Fig. 1A(b)) matches with that of the bulk, cubic inverse spinel Fe₃O₄ phase (Fig. 1A(a)) (JCPDS 75-449; ICSD-029129). It has a space group of *Fd* $\bar{3}m$ S (227) and a cell volume of 580.09 Å³.

It was inferred that the synthesized nanomagnetic material is Fe₃O₄ in the cubic inverse spinel structure. The substitutional doping of Mn²⁺ in the Fe₃O₄ lattice showed certain distinctive features as outlined in Table 1. The XRD patterns of [Mn-1 to Mn-5] doped samples did not show any characteristic signature of Mn₂O₃, any other oxide of Mn, or even metallic Mn.

However, elemental analysis by energy-dispersive X-ray fluorescence (EDXRF) (Fig. S1) showed the presence of Mn in the Mn-doped Fe₃O₄ samples, which proved that Mn was



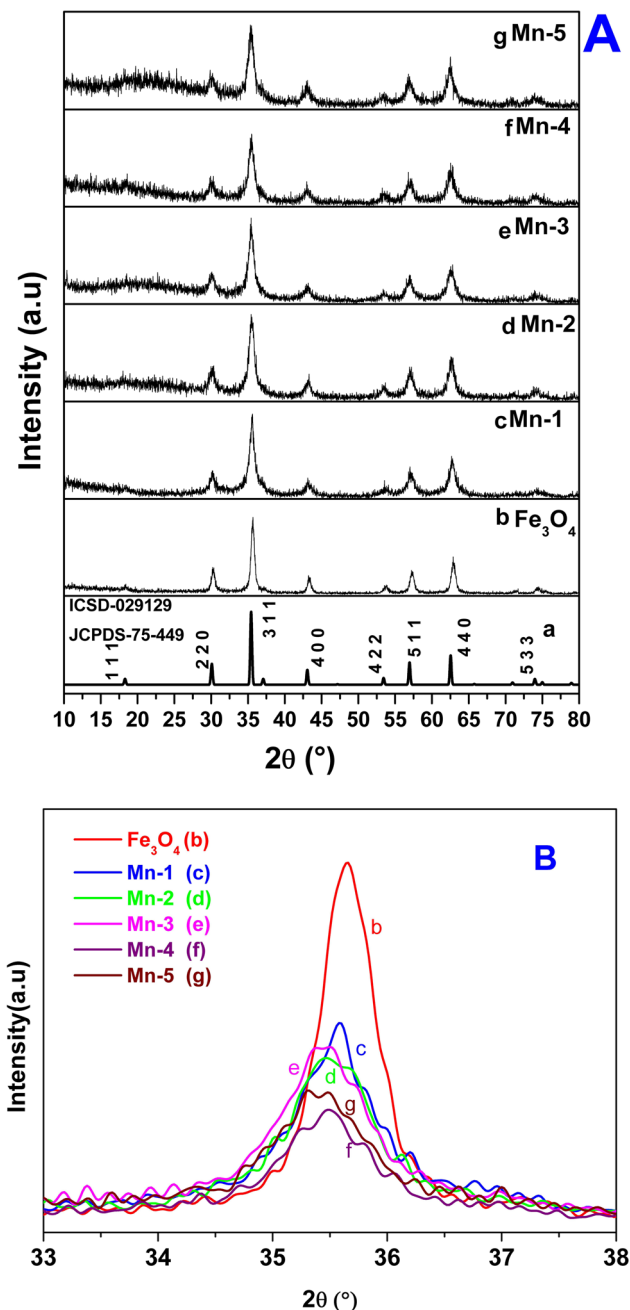


Fig. 1 (A) X-ray diffraction data for the synthesized MNPs and Mn-MNPs: (a) Fe₃O₄ of ICSD-029129, JCPDS75-449; (b) nano Fe₃O₄; (c) Mn-1; (d) Mn-2; (e) Mn-3; (f) Mn-4; (g) Mn-5. (B) Plots of the only (3 1 1) planes for the above samples: (a) nano Fe₃O₄, (b) Mn-1, (c) Mn-2, (d) Mn-3, (e) Mn-4, (f) Mn-5.

substitutionally doped in the cubic Fe₃O₄ lattice. The oxidation state of the doped Mn will be probed later in the XPS section. The Mn-MNPs (Mn-1 to Mn-5) (Fig. 1A(c)–(g)) showed the same phase as that of nano-Fe₃O₄. Although all the XRD patterns (for the Mn-doped samples) showed a single-phase character, the XRD lines of the nano Fe₃O₄ and Mn-doped Fe₃O₄ samples exhibited significant broadening as compared to the parent nano Fe₃O₄ along with a slight shift in the 2θ value of the

primary (3 1 1) plane as shown in the Fig. 1B, which showed strain due to the doping of Mn in the Fe₃O₄ lattice. The coherently diffracting domain sizes calculated using the Scherrer equation from the broadening of the FWHM of the (311) peak are shown in Table 1. The crystallite size of nano Fe₃O₄ was about 9.95 nm. The average crystallite size of the Mn-doped Fe₃O₄ [Mn-1 to Mn-5] materials was reduced as a function of the Mn²⁺ doped into the Fe₃O₄ lattice, and is also presented in Table 1. The crystallite size for Mn-1 is ~9.41 nm and that of Mn-5 is ~8.72 nm.

The ionic radius of Mn²⁺ is 83 pm (coordination number VI and high-spin state) and that of Fe²⁺ is 78 pm (coordination number VI and high-spin state).⁵⁷ Similarly, the covalent character of the Mn²⁺–O bond will be lower as compared to that of the Fe²⁺–O bond, therefore, the bond length of Mn–O is expected to be greater as compared to that of Fe–O.⁵⁸ The lattice parameters of the Mn-doped samples are included in Table 1. There is a substantial expansion in the Fe₃O₄ lattice parameters [*a* = *b* = *c*-axis (cubic) and the volume] as a function of the Mn²⁺ dopant concentration in the Fe₃O₄ lattice. Mostly, Mn²⁺ substitutes for the octahedral Fe²⁺ in the Fe₃O₄ lattice (as is shown in the subsequent sections) and, therefore, would cause an increase in the lattice parameters and volume of the lattice.

3.2 Raman spectroscopy

Fig. 2 shows the Raman spectra of the Mn-doped Fe₃O₄ samples including nano Fe₃O₄. Fe₃O₄ possesses an inverse spinel structure with a space group of *Fd3m* with a typical AB₂O₄ structure [A–Fe³⁺ & B–Fe²⁺/Fe³⁺]. The point group for the above Fe₃O₄ lattice is in the static configuration *D*_{3d} and if considered overall, it is octahedral (*O*_h). According to the factor group analysis, considering the *O*_h point group, there are 15 different vibrational modes of which 6 are Raman active, 6 are IR active, and 3 are photoacoustic modes.⁵⁹

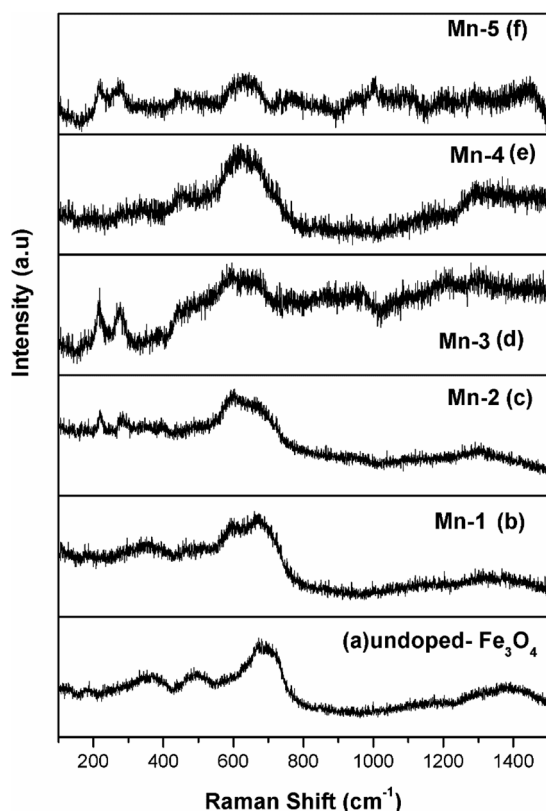
Fig. 2a shows three Raman peaks for nano Fe₃O₄: 671 cm^{−1} (A_{1g}), 510 cm^{−1} (T_{2g}), and 350 cm^{−1} (E_g). The other peaks were not observed for the present system. Mn-1 and Mn-2 (Fig. 2b and c.) also possess the same set of peaks as Fe₃O₄. Two new peaks (275 and 217) cm^{−1} were observed for the Mn-3 and Mn-5 (Fig. 2d and f). These peaks are due to γ-Fe₂O₃, which is known to form under the thermal oxidative effect of the laser during Raman spectroscopy.⁶⁰ No Raman peaks for the MnO_x were observed. This proves that Mn is doped in the Fe₃O₄ lattice.⁶¹ With greater doping of Mn in the Fe₃O₄ lattice, Raman peaks of Fe₃O₄ were observed, which invariably indicated that Mn was doped in the Fe₃O₄ lattice and there were no other segregated phases of any MnO_x in the Mn-doped Fe₃O₄ system.

3.2.1 BET surface area. Table 1 lists the specific surface areas of all the samples synthesized during the current study. All the samples synthesized by this modified co-precipitation route exhibited high surface areas in the range of 90–170 m² g^{−1}. Among them, Mn-4 had the highest surface area of 171 m² g^{−1}, followed by Mn-2, and Mn-1. The Mn-5 sample exhibited the lowest surface area of 129 m² g^{−1}. However, the surface areas of the Mn-doped samples are larger as compared to the nano Fe₃O₄ sample but the increment is not substantial.

Table 1 Textural, lattice and sorption parameters for different samples of MNPs and Mn-MNPs

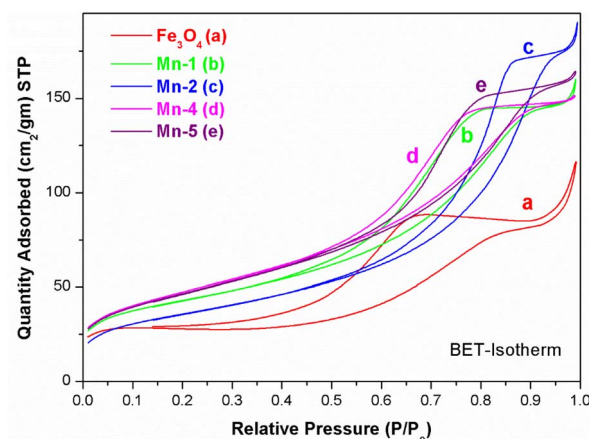
Sample	BET surface area ^a (m ² g ⁻¹)	Pore size ^b (Å)	Pore volume (cm ³) ^c	Type of isotherm	Crystallite size (nm) ^d	Unit cell parameter ($a = b = c$) ^e (Å)	Unit cell volume (Å ³)
Fe ₃ O ₄	89.03	60.52	0.19	IV	9.95	8.346(2)	581.54(0.28)
Mn-1	153.03	60.71	0.26	IV	9.41	8.369(4)	586.24(0.51)
Mn-2	168.54	80.80	0.30	IV	8.84	8.372(7)	586.77(0.88)
Mn-4	171.04	50.70	0.24	IV	8.77	8.383(9)	589.16(1.13)
Mn-5	129.14	56.47	0.26	IV	8.72	8.389(6)	590.59(0.83)

^a BET surface area from the linear portion of the BET plot in the relative pressure range of (P/P_0) 0.05–0.30. ^b Average pore diameter estimated using the BJH desorption branch of the isotherm. ^c BJH desorption cumulative volume of pores between 17.000 Å and 3000.000 Å in diameter. ^d Crystallite size measured from the Scherrer equation. ^e Unit cell parameters for the MNPs and Mn-MNPs are calculated using the POWDERX software, where Fe₃O₄ matches the JCPDS file (#75-0449); ICSD (#029129) showing the $Fd\bar{3}m$ (227) space group with the cubic unit cell structure of $a = b = c = 8.31$ Å, $\text{vol} = 580.09$ Å³ $\alpha = \beta = \gamma = 90$.

**Fig. 2** Raman spectra of the synthesized MNPs and Mn-MNPs: (a) nano Fe₃O₄; (b) Mn-1; (c) Mn-2; (d) Mn-3; (e) Mn-4, and (f) Mn-5.

The average pore size and total pore volume were also found to be almost similar for all the Mn-doped Fe₃O₄ samples (Table 1). Fig. 3 illustrates the typical nitrogen adsorption–desorption isotherms obtained for various Mn-doped Fe₃O₄ samples.

The type IV (IUPAC) hysteresis loop indicates the presence of mesoporous character in the sample. All the samples showed similar hystereses in their respective isotherm plots with inflection points in the P/P_0 range of 0.45–0.7, suggesting mesoporosity in these samples as presented in Table 1. The inflection seen at P/P_0 of ~0.9 may be attributed to the macropores caused by particle–particle porosity.

**Fig. 3** BET isotherms for the different samples of MNPs and Mn-MNPs: (a) Fe₃O₄; (b) Mn-1; (c) Mn-2; (d) Mn-4 and (e) Mn-5.

In Table 1, the units are represented in their generic terms, however, the S.I units for the above are as follows: 1 Å = 10⁻¹⁰ m; 1 nm = 10⁻⁹ m; 1 cm³ = 10⁻³ m³ kg⁻¹; 1 m² g⁻¹ = 10⁻³ m² kg⁻¹.

3.3 Electron microscopy studies

3.3.1 SEM and EDS studies. The Scanning Electron Microscopy (SEM) data for the samples are shown in Fig. 4 and their representative EDS data are shown in the corresponding lower half of the figure. The SEM studies showed typical spherical agglomerated structures of Fe₃O₄ (Fig. 4A(a)) and the EDS plot shows the presence of Fe in the system. Fig. 4A(b) presents the SEM features of Mn-1 showing similar features. The elemental analysis, in atomic weight percentage, by the EDS technique is presented in Table S1,[†] which reflects the increment in the Mn-percentage as a function of Mn-doping in the Fe₃O₄ lattice. Fig. 4B shows the elemental mapping for the Mn-4 sample; it shows the presence of all the elements in a mostly uniform fashion and the effective substitutional Mn doping is done uniformly.

3.3.2 TEM and HR-TEM studies. Representative transmission electron microscopy (TEM) and high-resolution transmission electron microscopy (HRTEM) images of nano Fe₃O₄



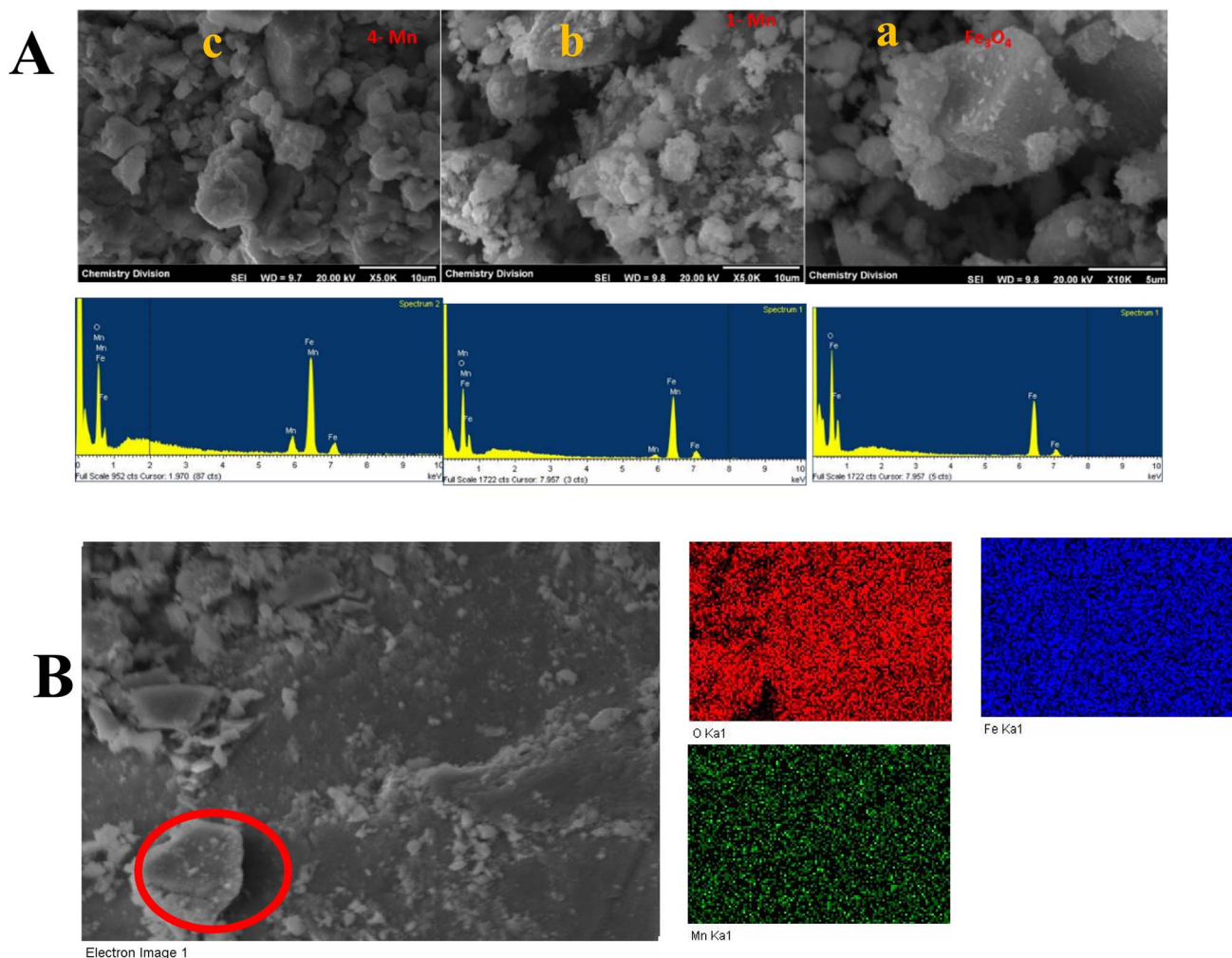


Fig. 4 (A) SEM images along with the EDS patterns of the synthesized (a) Fe₃O₄, (b) Mn-1, (c) Mn-4. (B) Elemental mapping of O, Fe, and Mn for Mn-4.

and Mn-3 samples are shown in Fig. 5A and C, respectively. TEM images of both samples show aggregates of nearly spherical particles (diameter 8–14 nm) as shown in (Fig. 5A(b) and C(d)). The HR-TEM and SAED (selected area electron diffraction) patterns shown in (Fig. 5A(a) and C(c)) revealed that these nanoparticles are highly crystalline. HRTEM images of both samples showed continuous lattice fringes, which suggests that the crystallinity is undisturbed by the doping of Mn in the Fe₃O₄ lattice. The distance between the lattice fringes was measured to be 3.028 Å, which corresponds to the (2 2 0) planes of nano Fe₃O₄ as shown in (Fig. 5B). Similarly, the distance between the lattice fringes of the Mn-3 sample is equal to 2.5765 Å, representing the (3 1 1) plane as shown in (Fig. 5D). Moreover, the diffraction rings in the corresponding SAED patterns are in good agreement with the XRD results discussed above (Tables S2† and 3).

3.4 Zeta potential

The zeta potential is majorly understood by the effective calculation of the surface potential of the formed double

Helmholtz layer over the surface for the Fe₃O₄ and Mn-doped Fe₃O₄ materials. The variation of the surface potential is an important factor in deciding the sorption of the uranyl ion (UO₂²⁺), which is a charged cation, over the surface of the Fe₃O₄ and Mn-doped Fe₃O₄ surface. Therefore, Fig. 6 shows the plot of zeta potential at different pH as a function of Mn-doping in the Fe₃O₄ system. It was observed that at lower pH (pH-3), the Mn-2% doped sample possessed the maximum positive zeta potential as a function of Mn-doping. However, at neutral pH, it was observed that upon doping with Mn, the surface zeta potential became more positive as shown in Fig. 6b, where the Mn-1 sample had the maximum positive zeta potential. Schemes 2 and 3 portray the hydration of the Mn-doped Fe₃O₄ system followed by the formation of the double Helmholtz layer to obtain a positive zeta potential as a function of Mn-doping in a neutral medium or at pH ~ 7.

The Mn-doped Fe₃O₄ would have mainly three types of cation, namely, Mn²⁺, Fe²⁺ and Fe³⁺ on its surface since M–O is present. As depicted in Scheme 2 this may be one of the many ways for the solvation of the Mn-doped Fe₃O₄. In this route,



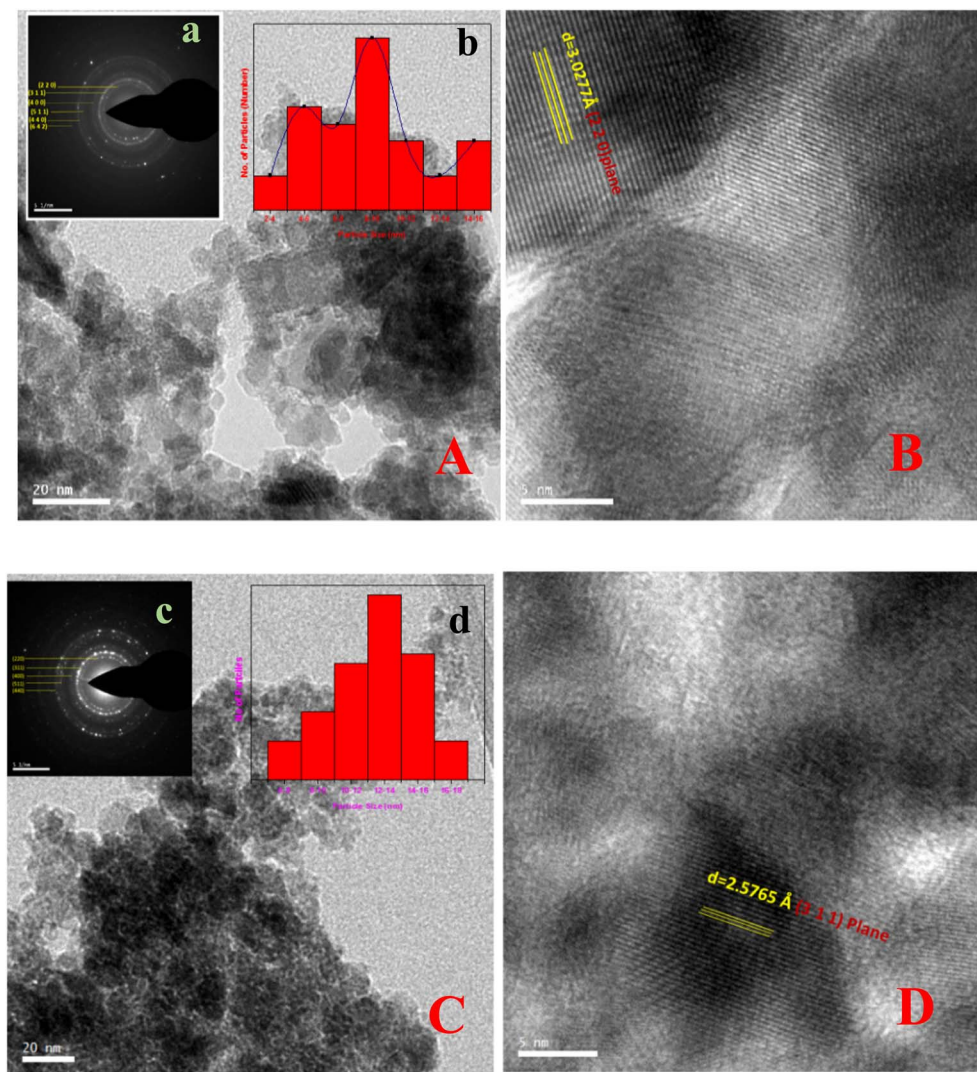


Fig. 5 TEM and HRTEM images along with the required d -spacings of Fe_3O_4 and Mn-3 samples: (A) (a) TEM image along with the SAED pattern (inset) with the assigned crystallographic plane, and (b) the particle size distribution of Fe_3O_4 particles (inset). (B) The HRTEM image with d -spacing and the required plane for Fe_3O_4 . (C) TEM images of Mn-doped Fe_3O_4 (Mn-3) along with the SAED pattern (inset) with the assigned crystallographic plane and particle size distribution of Mn-3 (inset). (D) HRTEM image with d -spacing and the required plane for Mn-3.

since M-O ($\text{M} = \text{Mn}^{2+}$, Fe^{2+} and Fe^{3+}) forms M-OH in the neutral medium, Mn^{2+} being the least electropositive among the three ions will attract less electron density from O in the described M-O bond and the O will therefore possess the maximum electron density [$\text{M}^{\delta+}-\text{O}^{\delta-}\cdots\text{H}-\text{O}-\text{H}$]; therefore, the formation of the outer stable hydration layer will be greatest for the Mn^{2+} . Clustering of the particles leads to an increase in particle size as observed in the TEM. The clustering of these particles accumulates more positive charge, leading to the formation of a higher positive surface charge and similarly higher zeta potential of Mn-doped Fe_3O_4 materials as observed in Fig. 6b.

As observed in Fig. 6a, in an acidic medium ($\text{pH} = 3$), the zeta potentials of the surfaces of all the Mn-doped Fe_3O_4 samples are mainly positive. It was also observed that an alteration in the zeta potential as a function of Mn-doping initially increased and had a maximum at the Mn-2 sample for Mn-doped Fe_3O_4 . The

formation of the double Helmholtz layer and its effect on the variation of the zeta potential in the acidic medium (*i.e.*, low pH) are shown in Scheme 4. The lower pH will possess excess H^+ ions, which will be adsorbed over the M-OH surface to form the A-type complex that will mainly have a surface with a positive charge, resulting in a positive surface charge. The Mn^{2+} will possess the least positive oxidation charge over it as compared to Fe^{2+} and Fe^{3+} . Therefore Mn^{2+} will attract a lower positive charge density from the surface layer and thereby, the surface will be more positive for Mn-doping as observed in Fig. 6a.

The zeta potential for the Mn-doped Fe_3O_4 initially increased and thereafter, the zeta potential for the higher-doped Mn^{2+} decreased as compared to Mn-1; this could mostly be due to the interparticle repulsion of the Mn^{2+} moieties that is bound to occur with an increase in the Mn^{2+} doping percentage in the Fe_3O_4 lattice.



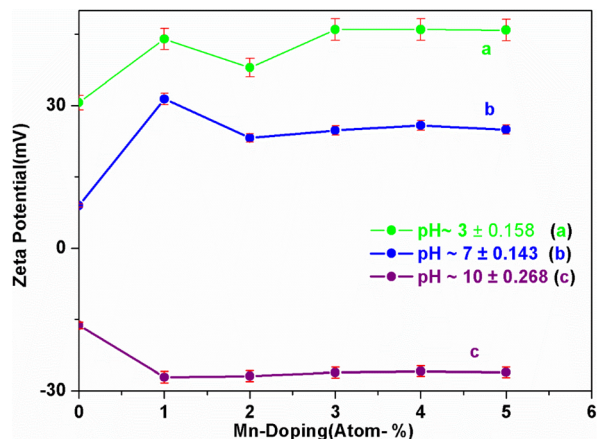
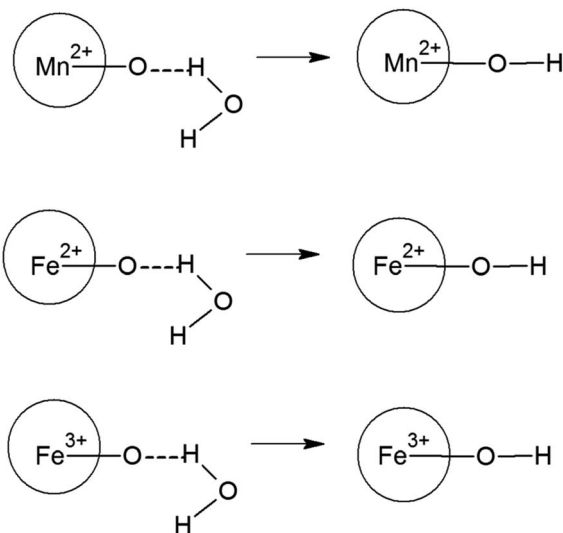
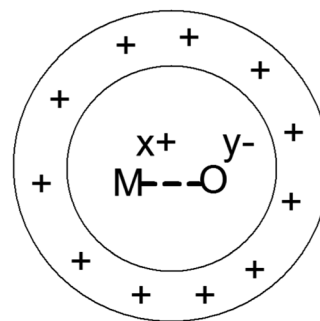


Fig. 6 Plots of zeta potentials at different pH: (a) pH-3; (b) pH-7 and (c) pH-10 as a function of Mn-doping in the Fe_3O_4 lattice. The bar at each point represents the error bar from the average of three sets of data taken for the same pH. This error along with the instrumental error contains the error in the pH calculation as shown in the legends. (Note: The S.I unit of zeta potential is V; $1 \text{ mV} = 10^{-3} \text{ V}$).

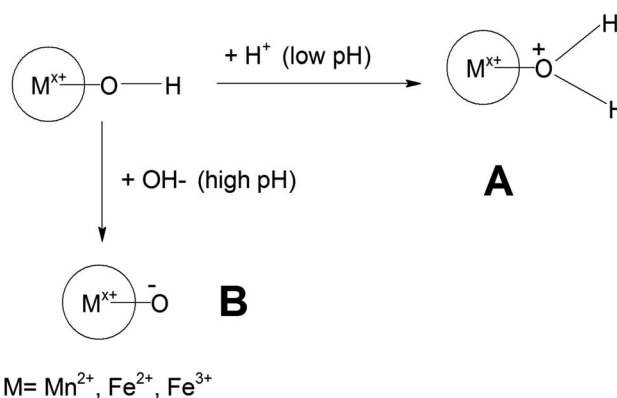


Scheme 2 Formation of the outer hydration layer with Mn-doped Fe_3O_4 . These schemes are mere representations and do not show the actual model.

Fig. 6c shows that the zeta potential for the Mn-doped Fe_3O_4 species is negative at a very high pH of ~ 10 . This could also be explained similarly by Scheme 4 as given above. In high pH media, there will be a high population of the OH^- ion. These OH^- ions will, in turn, extract an H^+ ion to form a complex ion like B representing the double Helmholtz layer. Therefore, the surface charge for this surface layer will be negative, as was observed in the experiments. With Mn-doping, the Mn will attract lower electron density from the surface as compared to that of Fe^{2+} and Fe^{3+} , thus making the surface charge of the Mn-doped Fe_3O_4 moieties more negative as compared to only Fe_3O_4 . This is also observed in Fig. 6c where the Mn^{2+} doping further lowers the zeta potential.



Scheme 3 Formation of double Helmholtz layer in the neutral medium. This scheme is a mere representation and do not show the actual model.



Scheme 4 Formation of the surface layer for the zeta potential in different media with low pH (acidic) and high pH (basic) in the present system.

3.5 XPS-studies

The XPS studies were conducted for the different Mn-doped Fe_3O_4 and nano Fe_3O_4 magnetic materials. Fig. 7A shows the Fe 2p XPS spectrum of the Mn-doped Fe_3O_4 along with that of nano Fe_3O_4 . Fe $2p_{3/2}$ and Fe $2p_{1/2}$ are shown at 710.8 and 724.8 eV, respectively, as has been reported for the Fe_3O_4 system in previous reports.^{62,63} In Fe_3O_4 , iron has two oxidation states, Fe^{2+} and Fe^{3+} , with the former being octahedrally coordinated and the latter distributed over both octahedral and tetrahedral sites to produce the inverse spinel structure.⁶⁴ These spectra were deconvoluted to understand the presence of the Fe^{2+} and Fe^{3+} present in the Fe_3O_4 system individually. Partially filled 3d levels result in the complex p-d mixing of the orbitals leading to the formation of the shake-up peaks,⁶⁵⁻⁶⁷ all of which contribute to the peak structure, leading to the formation of the satellite peak in Fe_3O_4 . However, in previous reports, satellite peaks were absent for the Fe_3O_4 system.⁶³⁻⁶⁹

For the Fe_3O_4 system synthesized by the co-precipitation method, we observed the presence of a satellite peak at 718.9 eV. The peak at 710.6 eV was attributed to Fe^{2+} and the peak at 712.3 eV is due to Fe^{3+} , which is consistent with

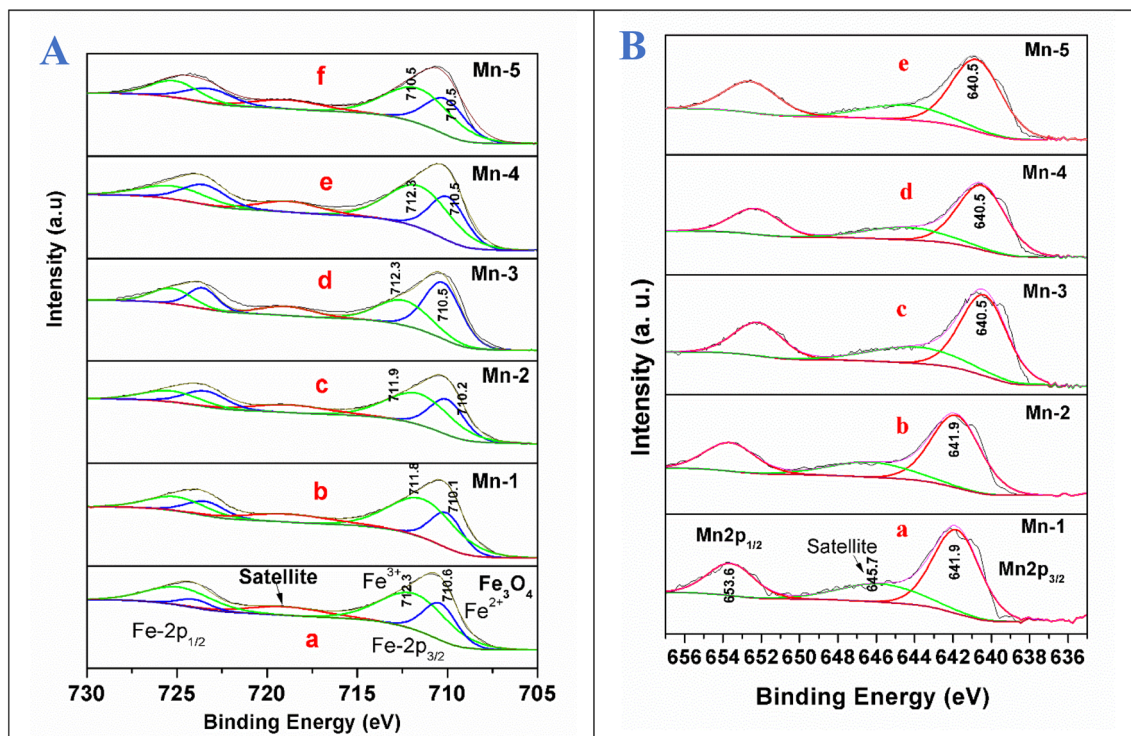


Fig. 7 (A) XPS spectra showing the deconvoluted Fe 2p ($2p_{3/2}$ and $2p_{1/2}$) peak for the samples: (a) nano Fe₃O₄; (b) Mn-1; (c) Mn-2; (d) Mn-3; (e) Mn-4; (f) Mn-5. (B) XPS spectra for the deconvoluted Mn 2p ($2p_{3/2}$ and $2p_{1/2}$), and the different nano Mn-doped Fe₃O₄ systems: (a) Mn-1; (b) Mn-2; (c) Mn-3; (d) Mn-4; (e) Mn-5. Note: It is worth mentioning that although the S.I. unit for energy is the Joule (J), in XPS terms, the binding energy (B.E.) is always represented in eV only ($1 \text{ eV} = 1.6021 \times 10^{-19} \text{ J}$). Therefore, in the entire text, B.E. is represented in eV units for brevity.

a previous report.⁶⁷ However, since the source of the present XPS is non-monochromatic Al K α , the further deconvolution of Fe³⁺ in the octahedral and tetrahedral states is not differentiable. With the substitutional doping of Mn in the Fe₃O₄ lattice, there was a very minor decrease in the binding energies of both Fe²⁺ and Fe³⁺ by 0.5 eV for Mn-1. In the other samples, the B.E. decreased to 0.4 eV for Mn-2, and 0.1 eV for the rest of the samples (Mn-3 to Mn-5) as shown in Fig. 7A. This shows that the electronic interactions between that of Fe³⁺/Fe²⁺ and that of the Mn ion were negligible, which mostly points towards the fact that Mn is doped as Mn²⁺ mostly replaces the Fe²⁺ sites, which is confirmed in a later section regarding the XPS results for the Mn 2p for the Mn-doped Fe₃O₄ systems.

Fig. 7B shows the Mn 2p XPS data for the different Mn-doped Fe₃O₄ samples. For the Mn-1 sample, the Mn 2p XPS spectrum shows the Mn 2p_{3/2} and Mn 2p_{1/2} peaks, respectively, at 641.9 and 653.6 eV, suggesting the presence of only Mn²⁺ as was observed previously.^{70–72} Upon the deconvolution of these peaks, only one peak could be fitted, showing the absence of any other oxidation state of Mn in the doped Mn in the Fe₃O₄ lattice. A satellite peak at 645.7 eV mostly shows the overlap with the valence d-band for Mn;^{73,74} in a transition metal, the satellite main peak is a function of several atomic parameters, of which the core overlap with the valence band is prime. The change in electronic density around Mn for the Mn-1 and Mn-2 samples was less, as shown in Fig. 7B(a). However, after Mn-2, there was a decrease in the binding energy by 1.4 eV from Mn-3 to Mn-5,

as shown in Fig. 7B(c–e). The doped Mn²⁺ is a d⁵ system, whereas the Fe²⁺ is a d⁶ system. Mn²⁺ mostly substitutes for the Fe²⁺ present in the octahedral positions of the Fe₃O₄. To fit the octahedral position, the Mn d⁵ system obtains more electron density to become stabilised in the octahedral sites only and not in other tetrahedral sites to form a disordered system. This is indirect evidence that shows that the Mn²⁺ mainly replaces the Fe²⁺-octahedral sites.⁷⁵

Fig. 8 shows the O 1s XPS spectra for the different Mn-doped Fe₃O₄ samples. The nano Fe₃O₄ possesses O 1s binding energy (B.E.) and a peak was observed at 529.8 eV, which can be deconvoluted into three peaks at 529.6, 530.3 and 531.9 eV, respectively (Fig. 8a). The B.E. corresponding to the first peak (529.6 eV) represents the lattice –O attached to Fe²⁺; the second peak (530.3 eV) corresponds to the lattice –O attached to Fe³⁺ and the third peak (531.9 eV) corresponds to the surface –OH groups present over Fe₃O₄. The Mn-1 sample has an O 1s peak at 529.8 eV and can be deconvoluted into four peaks at 529.2, 529.9, 530.7, and 532.2 eV, respectively (Fig. 8b). The first three O 1s peaks represent lattice O-atoms where the first O (529.2 eV) is the one attached to Mn²⁺, the next 529.9 eV represents that for O– attached to Fe²⁺ and the third one to that of Fe³⁺. The last O 1s represents the surface –OH groups for Mn-1-doped Fe₃O₄. Similarly, for the other Mn-doped Fe₃O₄ samples, *viz.* Mn-2, Mn-3, Mn-4 and Mn-5, the respective O 1s spectra are given in Fig. 8(c–f). As a function of Mn-doping, for Mn-3 to Mn-5, the binding energy for the O-attached to Mn increased from 529.2



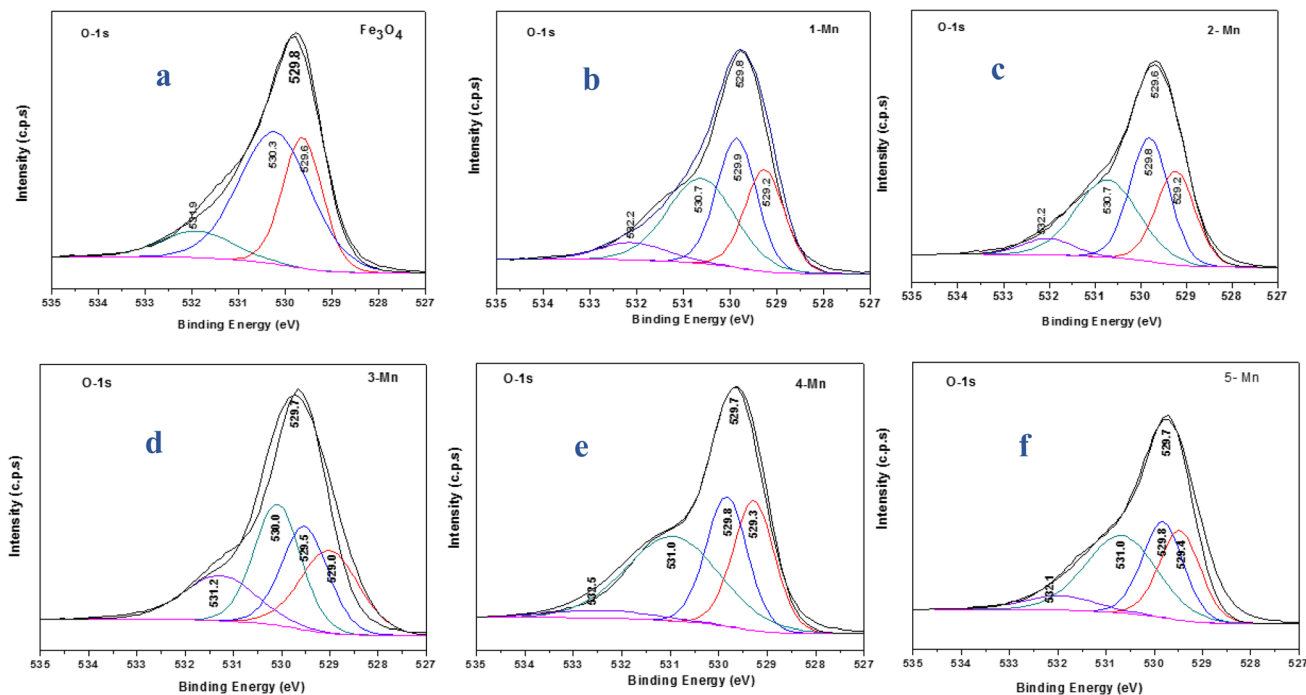


Fig. 8 XPS spectra showing the deconvoluted O 1s peaks for the different samples of MNPs and Mn-MNPs: (a) nano Fe_3O_4 ; (b) Mn-1; (c) Mn-2; (d) Mn-3; (e) Mn-4; (f) Mn-5.

to 529.4 eV, showing that Mn^{2+} mainly draws electron density from the $-\text{O}$ of the $\text{Mn}-\text{O}$ bond, thereby maintaining the $\text{Mn}-\text{O}$ bond length, the same as that of the $\text{Fe}-\text{O}$ ($\text{Fe}^{2+}-\text{O}$) octahedral bond length. However, the O 1s representing $\text{O}(\text{Fe}^{2+}-\text{O})$ and $\text{O}(\text{Fe}^{3+}-\text{O})$ remained almost unaltered throughout the series. Therefore, it can be deduced that the Mn^{2+} substitutes for the Fe^{2+} sites in the Fe_3O_4 inverse spinel lattice.

Fig. 9A shows the comparison of XPS data for Fe_3O_4 and Mn-doped Fe_3O_4 after uranyl ion adsorption over their respective surfaces. In Fe_3O_4 there is a significant lowering of the binding energy (1.2 eV) of Fe^{2+} after the uranyl ion adsorption over the Fe_3O_4 surface. This has been shown in Fe 2p XPS data portrayed in Fig. 9A(a and b).

It is quite apparent that Fe^{3+} remained almost unaltered even after the uranyl adsorption (711.8 to 711.2 eV, respectively). This is a clear sign that the uranyl ion is adsorbed mostly over the Fe^{2+} ions in the surface of Fe_3O_4 as compared to the Fe^{3+} ions, as shown in Scheme 5. This is mainly because the uranyl ion (UO_2^{2+}) is a cation with a charge of +2 and would have lower electronic repulsion with Fe^{2+} sites. Therefore, Fe^{2+} (d^6 electrons) possesses more electron density as compared to Fe^{3+} (d^5 electrons) in the Fe_3O_4 inverse spinel lattice. Upon adsorption, the electronic interaction is mostly with the O of uranyl ions and that of Fe^{2+} in Fe_3O_4 . Thus, upon adsorption of the uranyl ion, a certain amount of electron density is shifted from O (uranyl ion) to Fe^{2+} (Fe_3O_4). Therefore, the post-adsorption Fe^{2+} possesses a lower binding energy as compared to the Fe^{2+} of the parent Fe_3O_4 . However, in Fig. 9B, which shows the O 1s XPS spectra for Fe_3O_4 and Fe_3O_4 after the adsorption of uranyl ions, the shift in the binding energy is nominal (~ 0.4 eV), which indicates that the O of Fe_3O_4 does not have substantial

interactions with the U^{6+} of the uranyl ion as has been portrayed in Scheme 5. Therefore, in the Fe_3O_4 surface, the contribution of the Fe^{2+} ions will mainly affect the process of adsorption for the uranyl ions. Fig. 9C presents the XPS comparison data of Mn 2p for the Mn-2 sample, where it shows the effect of the uranyl ion adsorption over the Mn-doped Fe_3O_4 sample. There is a definite lowering in the binding energy for the Mn 2p (~ 1.7 eV) where the Mn 2p shifts from 641.9 to 640.2 eV after the uranyl ion adsorption. This also points towards a definite interaction between the substituted Mn^{2+} ion in the Mn-doped Fe_3O_4 lattice and that of the uranyl ion. The interaction between the Fe^{2+} or Fe^{3+} present in the Mn-doped Fe_3O_4 is shown in Fig. S2,† where it is reflected that there was very little alteration in the binding energy of the $\text{Fe}^{2+}/\text{Fe}^{3+}$ after uranyl ion adsorption over the Mn-doped Fe_3O_4 . Hence, it may be concluded that the uranyl ion mainly interacts with the Mn^{2+} ions of the Mn-doped Fe_3O_4 surface. This can also be understood from the similar explanation given above. The uranyl ion, being charged, will interact with the most electron-dense part of the surface, namely, the Mn^{2+} ions. This is reflected in Scheme 6. Fig. 9D presents a comparison of the O 1s XPS peaks for the same Mn-doped samples after the uranyl ion adsorption in the system. Similarly, the lowering of the binding energy for the O 1s system is negligible (~ 0.4 eV) and Scheme 6 shows that there is very little or negligible interaction between the U^{6+} of the uranyl ion and O in the Mn-doped Fe_3O_4 lattice surface.

3.6 Magnetic studies at room temperature

The VSM results for the magnetic study are shown in Fig. 10 for the different samples of MNPs and Mn-MNPs at room



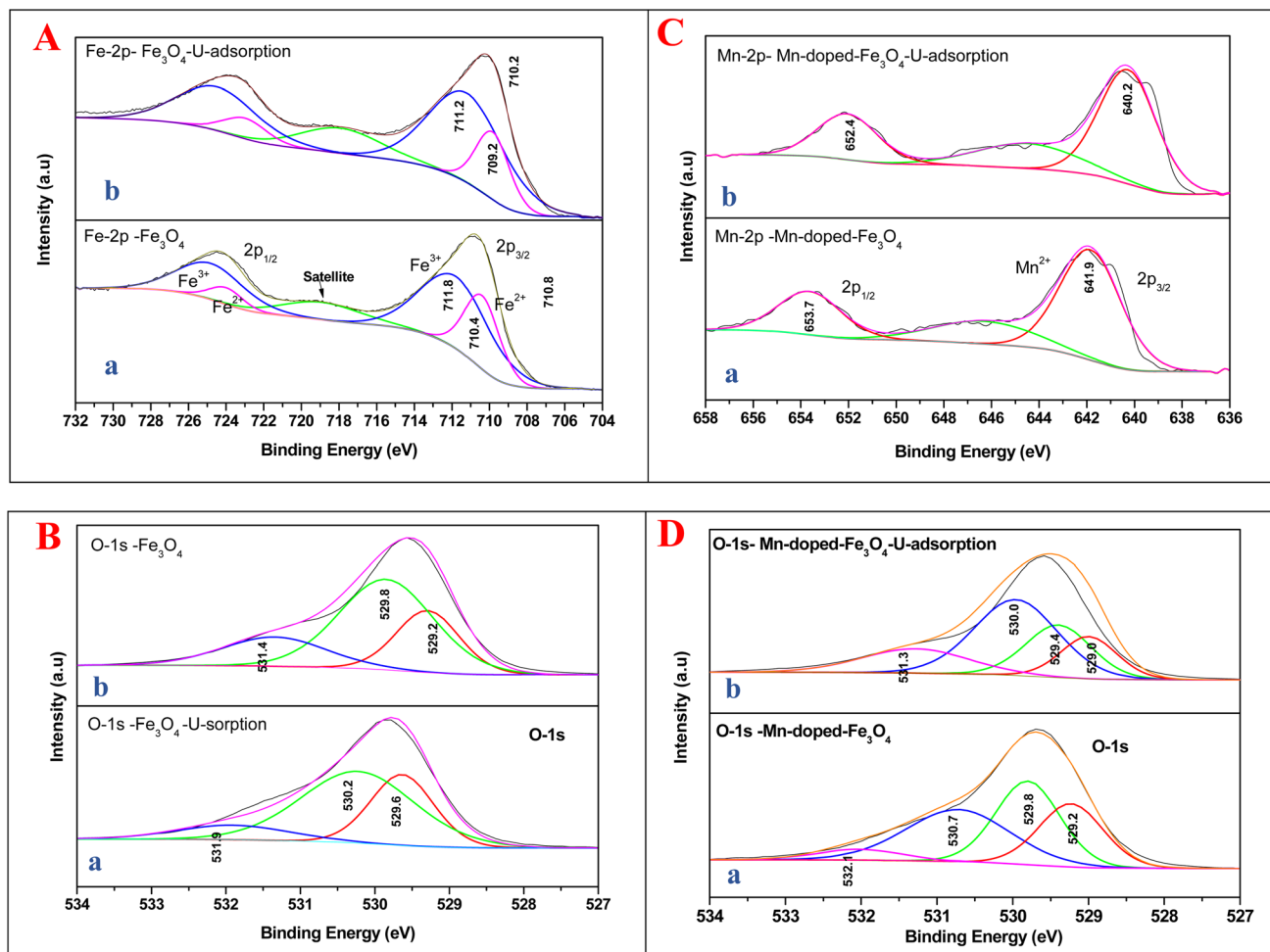
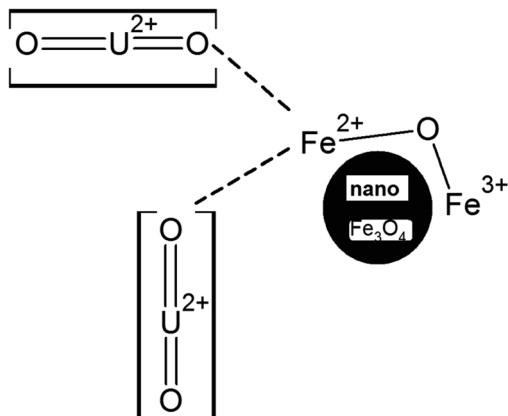


Fig. 9 XPS spectra for the comparison of the process of sorption interaction: (A) Fe 2p for (a) Fe₃O₄ and (b) Fe₃O₄ after U sorption post-drying. (B) O 1s – (a) Fe₃O₄ and (b) Fe₃O₄ after U sorption post-drying. (C) Mn 2p for (a) Mn-2 and (b) Mn-2 after U adsorption and post-drying. (D) O 1s for (a) Mn-2 and (b) Mn-2 after U adsorption and post-drying.

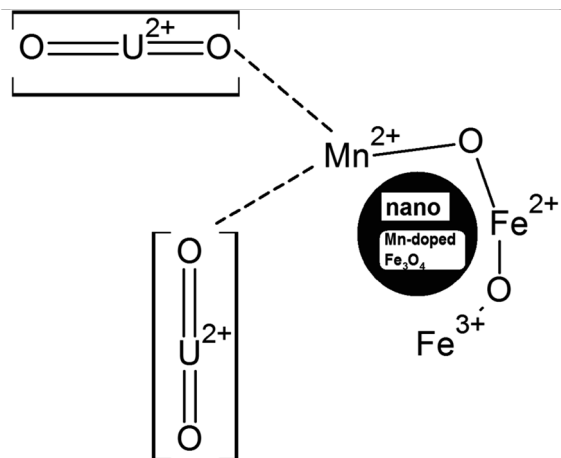


Scheme 5 Representation of the binding of Fe²⁺ sites with the UO₂²⁺ ions in the synthesized Fe₃O₄ nanoparticles. The above schematic representation is not indicative of molecular-level bonding phenomena.

temperature only. The results show the magnetic moments for all the Mn-MNPs to be greater as compared to the Fe₃O₄ samples.

The above room temperature VSM studies show the magnetic moments for the samples as 36.74, 42.44, 55.2, 44.4 and 45.8 emu g⁻¹ [(S.I unit for magnetization is A m⁻¹); 1 emu g⁻¹ ≡ 5.17 × 10³ A m⁻¹ for Fe₃O₄ & Mn-doped Fe₃O₄ samples; the dopant percentage is negligible so the density for all the materials can be considered the same as Fe₃O₄] for the samples Fe₃O₄, Mn-1, Mn-2, Mn-4 and Mn-5, respectively. The magnetic moment initially increased and at a greater concentration of the Mn-dopant in the Fe₃O₄ lattice, the magnetic moment decreased. It is quite evident from the XPS data that in Mn-doped Fe₃O₄, the Fe²⁺ sites are substituted by the Mn²⁺ ions. Mn²⁺ is a d⁵ system with a higher magnetic moment, whereas the Fe²⁺d⁶ system possesses a lower magnetic moment as compared to Mn²⁺. This phenomenon is manifested up to the Mn-2 sample where with a higher concentration of Mn²⁺ dopant, the magnetic moment increases as observed in Fig. 10. This can also be an alternate proof showing that for the Mn-





Scheme 6 Representation of the binding of Mn^{2+} sites with the UO_2^{2+} ions in the synthesized Mn-doped Fe_3O_4 nanoparticles. The above schematic representation is not indicative of molecular-level bonding phenomena.

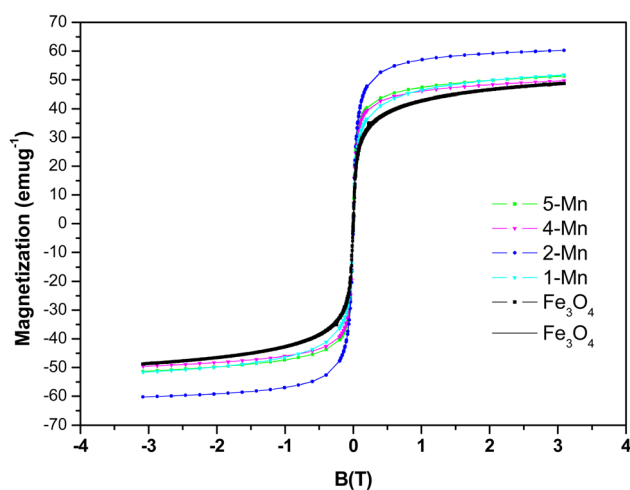


Fig. 10 Magnetization of the different Mn-doped samples over Fe_3O_4 .

doping in the Fe_3O_4 lattice, the Fe^{2+} sites are mainly substituted by the Mn^{2+} ions.

However, it has been observed by Warner *et al.*,⁴⁹ that a higher concentration of Mn^{2+} doping (>5 wt%) resulted in lowering magnetic moments. This has been ascribed to the transition from magnetite to maghemite with its lower saturation magnetization (M_s), resulting in the distribution of Fe^{3+} ions on the tetrahedral and octahedral sites to a greater extent as compared to the Fe_3O_4 lattice. In addition, previous literature also shows a strong dependence of M_s on the particle size, which has similarly been stated earlier for the Mn-doped Fe_3O_4 systems. Here, these nanoparticles experience spin canting over the surface, thereby lowering their magnetic moment, which is probably true for these Mn-MNPs.⁷⁶ Table 1 shows that as a function of Mn-doping, the crystallite decreases, which will further augment the spin canting term to a greater extent in the highly Mn-doped Fe_3O_4 systems.

3.7 Sorption studies for uranyl ions

3.7.1 Acidic medium (pH 3). The adsorption of uranyl ions taken as $\text{U}(\text{nat.})$ standard (~ 75 ppb) was utilised to understand the sorption of the uranyl ions over the MNPs and Mn-MNPs at different pH as a function of Mn doping and was analysed using a LED fluorimeter. Fig. 11A shows the sorption of the U-ion over nano MNPs and Mn-MNPs samples in an acidic medium ($\sim \text{pH } 3$). It was observed that all the Mn-doped Fe_3O_4 samples were better sorbing agents.

Almost all the sorbents including the MNPs and the Mn-MNPs adsorbed nearly 100 percent of the uranyl ions within ~ 60 minutes. Fig. 11 shows the good sigmoidal fitting of the sorption parameters, indicating that the sorption behaviors of all the sorbents followed first-order kinetics. Fig. 11A shows that the Mn-2 sample followed the fastest kinetics for sorption. To quantitatively describe the kinetics of the sorption, the $t_{1/2}$ of the sorption rate was determined, where " $t_{1/2}$ " is calculated as the time in which 50 percent of the uranyl ions are adsorbed over the MNPs and the Mn-MNPs. This is taken as a kinetic

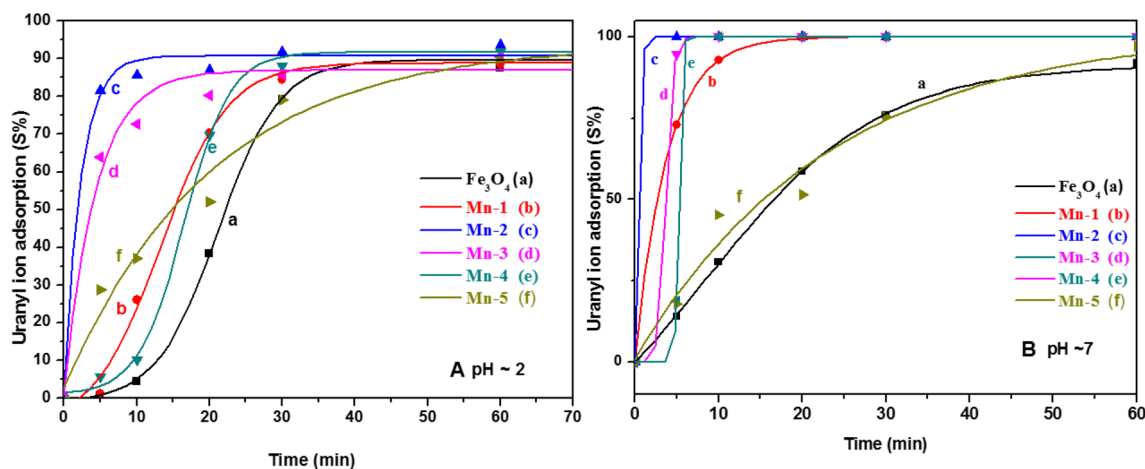


Fig. 11 The adsorption of uranyl ions (S%) on different samples of MNPs and Mn-MNPs at (A) $\text{pH } 3$ and (B) $\text{pH } \sim 7$ for (a) nano Fe_3O_4 ; (b) Mn-1; (c) Mn-2; (d) Mn-3; (e) Mn-4; (f) Mn-5.



factor to understand the efficiency of the kinetics of the different MNPs.

Adsorption is also represented by distribution coefficient (K_d) (eqn 1) values that were calculated from the analyzed post-contact solution and reference solution activity values using the following equation: (eqn 3)

$$K_d = \frac{C_i - C_f}{C_f} \times \frac{V}{M} \quad (1)$$

where C_f = the final concentration (ppb); C_i = initial concentration (ppb); V = volume of sample solution (mL); M = weight of sorbent taken (g).

Sorption percentage ($S\%$) (eqn 2) as in Fig. 11, plotted with time, can be defined as

$$S(\%) = \frac{C_i - C_f}{C_f} \times 100 \quad (2)$$

The $t_{1/2}$ values and the K_d values for the sorption reactions over the MNPs and the Mn-MNPs are plotted in Fig. 12. With the lowest $t_{1/2}$ and highest K_d for the sorption of uranyl ions over the Mn-2 sample (Fig. 12) in the acidic and neutral medium, Mn-2 is the best material among the MNPs and Mn-MNPs synthesized in this study for uranyl ion sorption. The $t_{1/2}$ values at different pH for the sorption of uranyl ions over the surface of MNPs and Mn-MNPs are well reflected in Fig. 12A(a). Fig. 12A(b) shows the different K_d values for the uranyl ion adsorption by the MNPs and Mn-MNPs. The K_d value is the maximum for Mn-2 in the neutral medium (Fig. S3†) and is almost equal to 3.08×10^6 , which is one of the best for uranyl ion sorption.

Two structural factors play a decisive role in the adsorption of the uranyl ions over the surfaces of these magnetic materials. Initially, from the XPS results described above, the generic model presented in Scheme 6 shows that in the Mn-doped Fe_3O_4 system, the uranyl ions are primarily adsorbed over the Mn^{2+} sites. Also, there are no defect sites over the surface of the

Mn-doped Fe_3O_4 sorbing materials, which could play an ionic role in the sorption of the positively charged uranyl ions over the Mn-MNPs surfaces. The XPS studies showed that there were more effective interactions of the sorption over Mn^{2+} as compared to the Fe^{2+} ions. Therefore, in the Mn-doped Fe_3O_4 system, the uranyl ions mainly interact with the Mn^{2+} system and are adsorbed over these sites.

The probability of sorption of the uranyl ions over the surface of the Mn-MNPs increases with an increase in at% of Mn^{2+} in the system. As the sorption of the doubly charged cation occurs over the surface, the surface charge of the sorbent will also play a strong role in the ionic factors for the surface. However, the surface charge as observed from the zeta potential data shows that it varies as a function of pH and thereby, will play different roles in different solvents. At an acidic pH, the surface charge was observed as positive and has been explained earlier in Section 3.4 in Scheme 4. The positive surface charge will mainly repel the positively charged sorbent over the surface. The positive surface charge for the Mn-MNPs samples initially decreases and is the lowest for Mn-2. Therefore, for Mn-2 the repulsion will be the least, making it the best sorbent for the uranyl ions at acidic pH. However, as a function of the Mn-doping, the surface charge (zeta potential) becomes more positive as a function of Mn-doping. This will be a detrimental factor for the sorption of the cations over these surfaces as is observed in Fig. 12. This is explained in Scheme 7.

3.7.2 Neutral medium (pH 7). Fig. 11B shows the trend for the sorption over the surface of the MNPs and the Mn-MNPs at neutral pH (pH 7). This trend is different from that at acidic pH and is explained below. The $t_{1/2}$ values for the MNPs and the Mn-MNPs are plotted in Fig. 12A(b). However, the representative K_d values for the same materials are presented in Fig. S3† since they were in a very high scale range and it was not possible to accommodate them in a single figure. At neutral pH, the best sorbent is Mn-2, however, the rates of sorption for all the MNPs

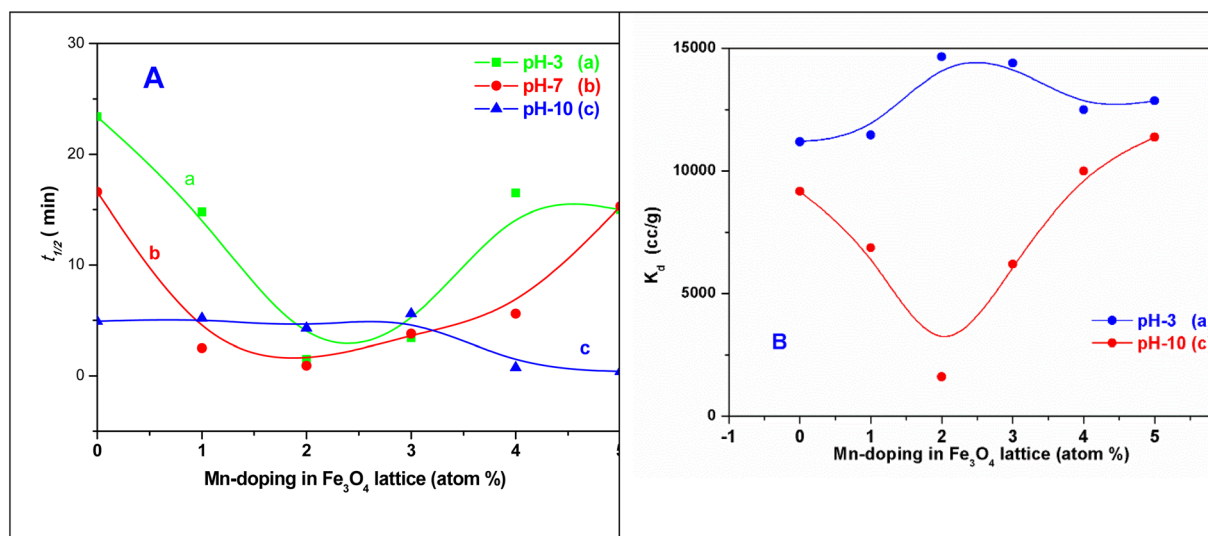
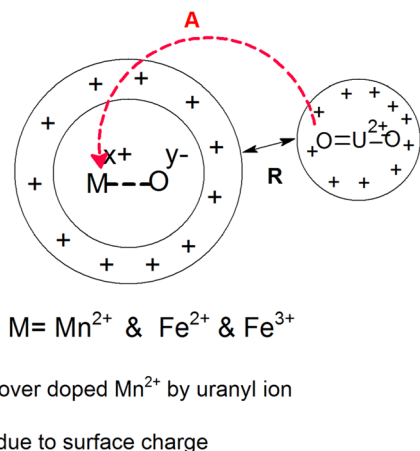


Fig. 12 Plots of (A) $t_{1/2}$ versus the Mn-doping in the Fe_3O_4 lattice in at% at (a) pH 3, (b) pH 7, and (c) pH 10 and (B) K_d versus the Mn-doping in the Fe_3O_4 lattice in at% at (a) pH 3 and (c) pH 10 K_d [pH 7 in the ESI†]. Note: SI unit for K_d L kg^{-1} ($1 \text{ cm}^3 = 10^{-3} \text{ L kg}^{-1}$).





Scheme 7 Ionic interactions for the sorption of uranyl ions over Mn-MNPs at low and neutral pH.

and Mn-MNPs are much as compared to acidic pH. This behaviour in the neutral medium can be explained by the above model presented in Scheme 7. Accordingly, the initial interaction of the Mn^{2+} and that of the uranyl ions will be the same, and constant in both the neutral and acidic media. However, the surface zeta-potential in the neutral pH is not the same as that of the acidic pH as observed in Fig. 6. The surface zeta potential is less positive as compared to that in the acidic medium. Therefore, the factor of repulsion from the surface positive charge will be much lower as compared to that of the acidic medium for the sorption of the positively charged cations of uranyl ions over the Mn-doped Fe_3O_4 samples. Therefore, the sorption kinetics as represented by both $t_{1/2}$ and K_d values can be well explained by the above hypothesis. Therefore, due to the least repulsion by the surface charge, the sorption in the neutral medium is the best and almost 100% of the uranyl ions are adsorbed within five mins.

3.7.3 Basic medium (pH 10). The sorption kinetics in basic media are presented in Fig. 13 for the MNPs and the Mn-MNPs. In a basic medium, the sorption of the uranyl ions over the different magnetic particles again tends to become slow and ~80% of the uranyl ions are completely sorbed over the surface. The trend in the sorption kinetics can be understood from the representative $t_{1/2}$ and the K_d parameters of the samples as plotted in Fig. 12A(c) and B(b). Here, the kinetics initially decreases for up to Mn-2 and then the sorption kinetics improves considerably and is almost constant for Mn-3 to Mn-5, which is reflected in both the $t_{1/2}$ and K_d values. In a basic medium, according to the literature,⁷⁷ the uranyl ions are mainly present as complex anions like $[\text{UO}_2(\text{OH})_3(\text{H}_2\text{O})_3]^{-1}$. The complexity of these ions varies and depends on the pH of the medium as shown in Fig. S5,[†] which shows the different species of the uranyl ion in the presence of nitrates at the required concentrations. Fig. 6c shows that the surface zeta potentials for the Mn-doped Fe_3O_4 nanoparticles are negative, where Mn-2 has the most negative zeta potential.

Therefore, the sorption of the uranyl ions can be understood from the model presented in Scheme 8. The interactions of the

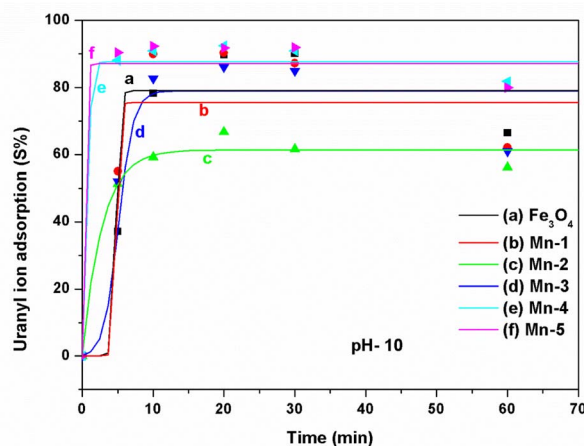
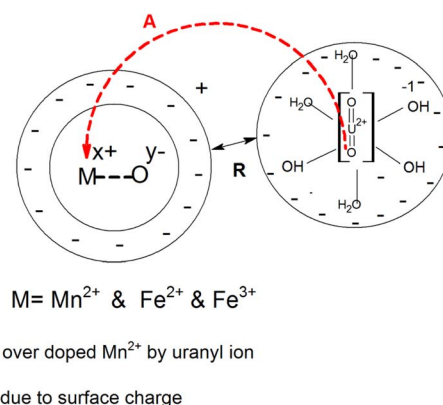


Fig. 13 The adsorption of uranyl ions (S%) on different samples of MNPs and Mn-MNPs in basic medium (pH = 10) for (a) nano Fe_3O_4 ; (b) Mn-1; (c) Mn-2; (d) Mn-3; (e) Mn-4; (f) Mn-5.

Mn^{2+} ions as a function of doping in the Fe_3O_4 surface with the uranyl ion will also be similar because they are in acidic or neutral media. However, because the surface zeta potential becomes negative and the uranyl ions in the basic medium also possess a negative charge, the repulsion between these becomes stronger. Similarly, the overall radii for the uranyl ion complexes will be larger in a basic medium as compared to that in an acidic or neutral medium. This will also result in further steric/ionic repulsion between the intermolecular uranyl ions, which will cause a further lowering of the adsorption kinetics in the basic medium.

3.8 Sorption studies for competitive ions

Since the study is about the sorption of uranyl ions from urine, studies of the sorption of certain inorganic ions generally present in urine along with uranyl ions were performed under simulative conditions. The ions studied were Na^+ , K^+ , Ca^{2+} and Mg^{2+} . As was observed in an earlier section, Fe_3O_4 and the pH of the medium play strong roles in altering the sorption properties



Scheme 8 Ionic interactions for the sorption of uranyl ions over Mn-MNPs at basic pH.



of these ions and, therefore, they were studied at acidic pH (\sim pH 4) and basic pH (\sim pH 9). The sorption of the ions is portrayed in Fig. 14 for three different MNPs and Mn-MNPs surfaces, *viz.* Fe₃O₄, Mn-2, and Mn-4, respectively.

The effects of the sorption of these competing ions in the nano Fe₃O₄ surface in either medium (pH \sim 4 & pH \sim 9) are negligible. At an acidic pH in the Mn-2 system, the sorption of these ions was found to be negligible but at basic pH, K⁺, Ca²⁺ and Mg²⁺ ions were adsorbed to the extent of \sim 25%. Competing ions like K⁺, Ca²⁺ and Mg²⁺ would presumably interact directly with either the Fe²⁺ in Fe₃O₄ or Mn²⁺ in Mn-doped Fe₃O₄ as observed in Schemes 5 and 6. Therefore, the interaction would be lower as compared to that of the uranyl ions because there, the Mn²⁺/Fe²⁺ interacts with the O of the uranyl ion whereas, here, it has direct interaction with either the solvated or the bare ions with positive charges, leading to less interaction, which explains the lower sorption of all these competing ions over these MNPs and Mn-MNPs surfaces. However, the surface charge also plays a significant role in the process of adsorption, which again can be explained by the previous model as proposed in Schemes 7 and 8. The Mn-2 system has a positive zeta potential at acidic pH and a negative zeta potential at basic pH. Therefore, there will be further repulsion for these competing ions at acidic pH, which will make the sorption negligible. However, in basic pH, since the surface is mostly negatively charged, there will be some attraction between these competing ions and the surface, leading to better sorption of the competing ions as shown in Fig. 14d. Similarly, for the Mn-4 system, the sorption is negligible at acidic pH (Fig. 14e) and there is considerable sorption of Ca²⁺ at basic pH, which can

also be explained as above. As shown by Table 1, there was not much alteration of the textural properties of the MNPs and the Mn-MNPs to substantially modify the sorption efficiency.

3.9 *Ab initio* calculations

3.9.1 Doping of Mn in the Fe₃O₄ lattice and the (3 1 1) surface of Fe₃O₄ and Mn-doped Fe₃O₄. The bulk structure of Fe₃O₄ was calculated from the coordinates obtained from the ICSD-029129 file. This ICSD file matches the experimental XRD data (Fig. 1a). The optimisation of the *K*-points for the Fe₃O₄ bulk (Table S 4†) shows K-3 3 3 to be the most optimised. The relativistic operations lead to the formation energy of -2.5530 eV per atom.

$$\Delta E_F = \frac{E_{\text{total}} - (n \times E_{\text{atom}})}{n} \quad (3)$$

where, ΔE_F = formation energy, E_{total} = total energy of the system, n = total number of atoms, E_{atom} = energy of individual atoms. Note: The S.I unit of energy is J; (1 eV = 1.6021×10^{-19} J). In the *Ab initio* calculations, ΔE_F eV per atom; surface energy eV \AA^{-2} [= $1.6021 \times 10^{-19}/10^{-20}$ J m^{-2} = 16.021 J m^{-2}]; slab energy eV, *etc.* The units are presented in eV units for brevity.

Mn was doped in the tetrahedral and octahedral sites of the Fe₃O₄ as shown in Fig. 15. Table S5† shows a comparison of the different lattice parameters of the Fe₃O₄ and Mn-doped Fe₃O₄ with the experimental results. Although the values for the *ab initio* calculations are not the same as the experimental values obtained from the Powderx software, the difference is not very great. Also, for the Mn-doped Fe₃O₄ (octahedral structural isomer) the trends in the lattice parameters and the cell volume

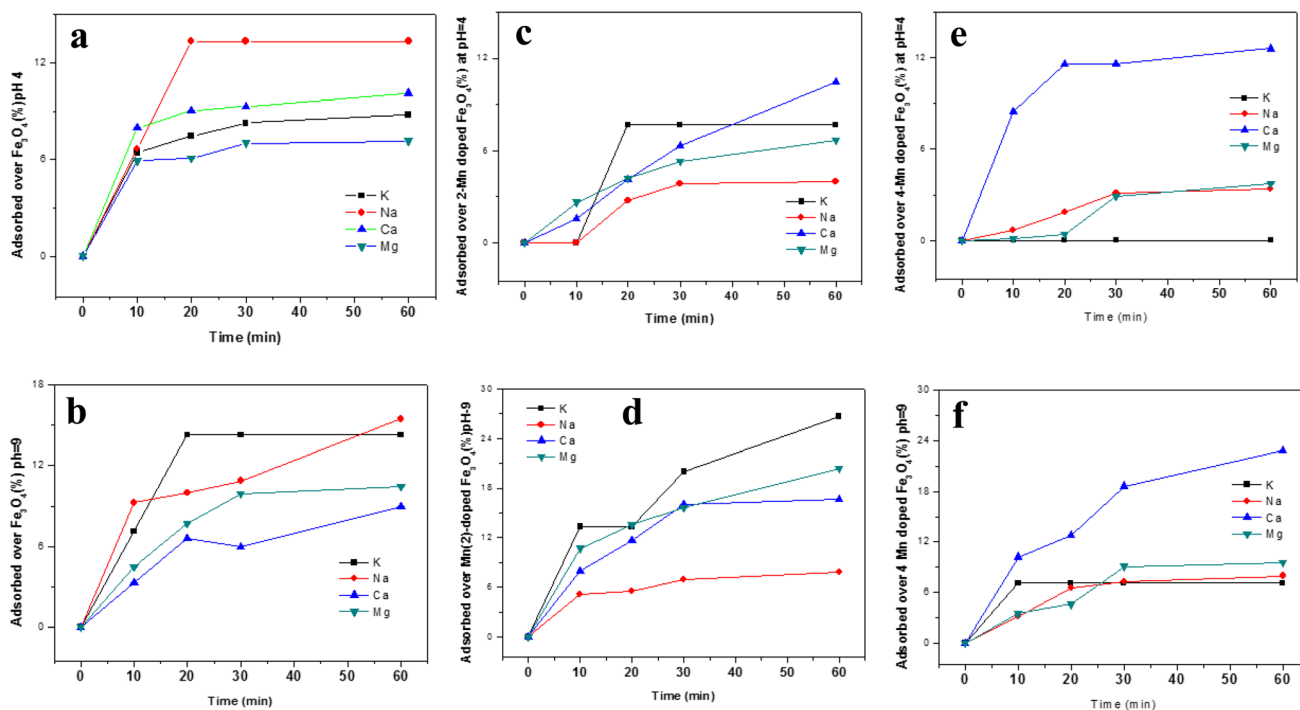


Fig. 14 The sorption of different ions (Na⁺, K⁺, Ca²⁺, Mg²⁺) over MNPs and Mn-MNPs surfaces: (a) nano Fe₃O₄ at pH 4; (b) nano Fe₃O₄ at pH 9; (c) Mn-2 at pH 4; (d) Mn-2 at pH 9; (e) Mn-4 at pH 4; (f) Mn-4 at pH 9.



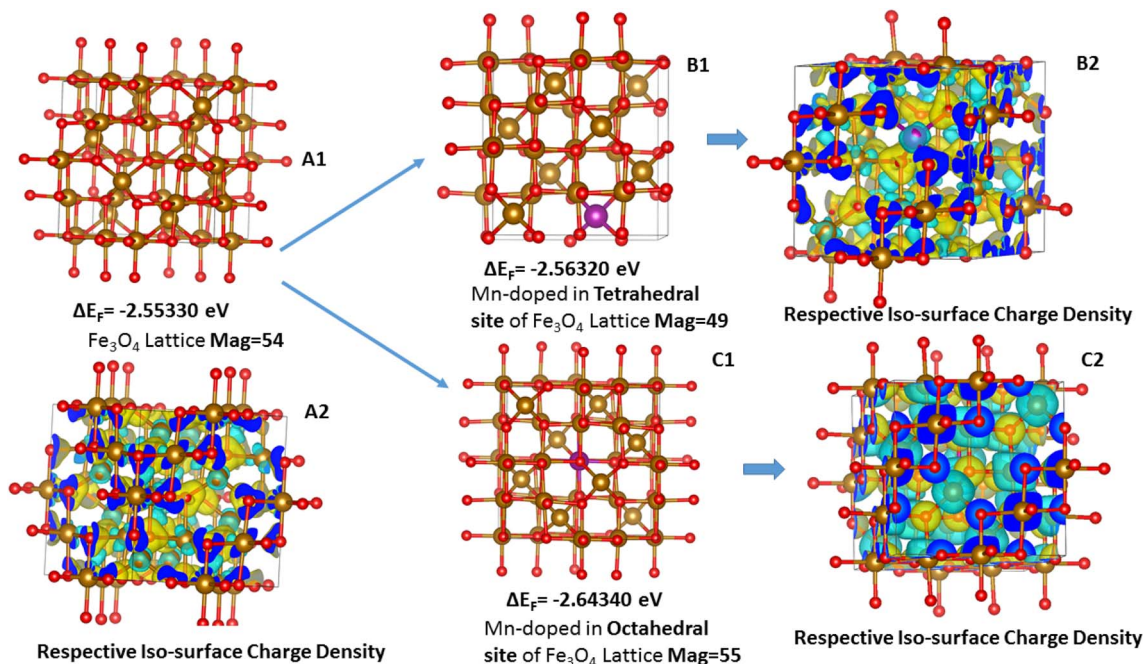


Fig. 15 Geometrical isomers for the doping of Mn in the Fe_3O_4 lattice: (A1) Fe_3O_4 lattice, (B1) Mn substituting the tetrahedral Fe-site; (C1) Mn substituting the octahedral Fe-site; (A2), (B2) and (C2) represent the charge density iso-surface plots of the A1, B1 and C1, respectively (eqn 3).

are the same as compared to the experimental value. The magnetic moment, though not calculated rigorously, also increases with Mn-doping as observed in Fig. 10 and is consistent with the experimental result. The geometrical isomer with octahedral substitution is the more stable isomer and the Octahedral Site Stabilization Energy per atom (OSSE) = $[-2.64340 - 2.56320] \text{ eV} = 0.08 \text{ eV}$, which reflects the fact that the doped Mn will preferably substitute for the Fe at the octahedral site as compared to that of the tetrahedral site. The relative charge that has been obtained from the XPS studies here shows that the Mn possesses a +2 charge as compared to a +3 charge. Therefore, it can be said that the Mn^{2+} will mainly substitute for the Fe^{2+} octahedral sites, which is reflected in the isosurface charge density.

The charge density potential iso-surfaces (Fig. 15) of Fe_3O_4 show that the two types of Fe in Fe tetrahedral sites and the Fe-octahedral sites have different depleted electron densities over the Fe atoms and electron accumulated densities over the O-atoms. The depleted electron density over the octahedral site (Fig. 15(C2)) is less as compared to that of the tetrahedral site (Fig. 15(B2)), which mostly shows that the Fe^{2+} is present at octahedral sites and Fe^{3+} is present at tetrahedral sites. The Mn-doping over the octahedral site shows that the Mn also possesses a similar electron-depleted density to the Fe octahedral site and, therefore, substitutes the Fe^{2+} sites as explained

earlier from the XPS results. Therefore, the octahedral Mn-doped Fe_3O_4 is the stable isomer (Table 2), which correlates with all the experimental parameters.

A (3 1 1) surface was cut both from Fe_3O_4 bulk and Mn-doped (Oct) Fe_3O_4 with 112 atoms and O on the final surface and was two monolayers thick. The (3 1 1) plane was taken because it is the major plane present in the XRD plots (Fig. 1b and c) as reported previously.⁷⁸ The surface created from the (3 1 1) plane was drawn in the form of a 2-layer slab model. Therefore, there were two layers of Fe_3O_4 (Fig. 16(A1)) and Mn-doped Fe_3O_4 (Fig. 16(B1)) with a total of 112 atoms.

$$\text{Surface Energy (S.E.)} = \frac{\text{slab energy} - (n \times \text{bulk energy})}{2 \times \text{area of (311) plane}} \quad (4)$$

The adsorption over the Mn-doped Fe_3O_4 surface is greater as compared to that of the Fe_3O_4 surface because the former is more stable, which also corroborates our experimental results and the XPS results. The charge density iso-surface plots for the (3 1 1) surfaces (Fig. 16(A2) and (B2)) for both Fe_3O_4 and Mn-doped Fe_3O_4 show that Mn is present as Mn^{2+} and replaces Fe^{2+} . Also, there is dispersed electron density over Fe and Mn along with that over O-atoms on the surface, which helps them to adsorb different molecules, including $[\text{UO}_2]^{2+}$.

Table 2 Surface energy of the (3 1 1) (eqn 4) surface of Fe_3O_4 and Mn-doped Fe_3O_4

Lattice system	Unit cell parameter a (Å)	Area of (3 1 1) surface (Å ²)	Slab energy (3 1 1) (eV)	Bulk energy (K-333) (eV)	Surface energy (eV Å ⁻²)
Fe_3O_4	8.43036	8.88638	−814.74718	−422.86294	1.743043
Mn-doped (Oct) Fe_3O_4	8.43887	8.89524	−824.80301	−426.99253	1.640318



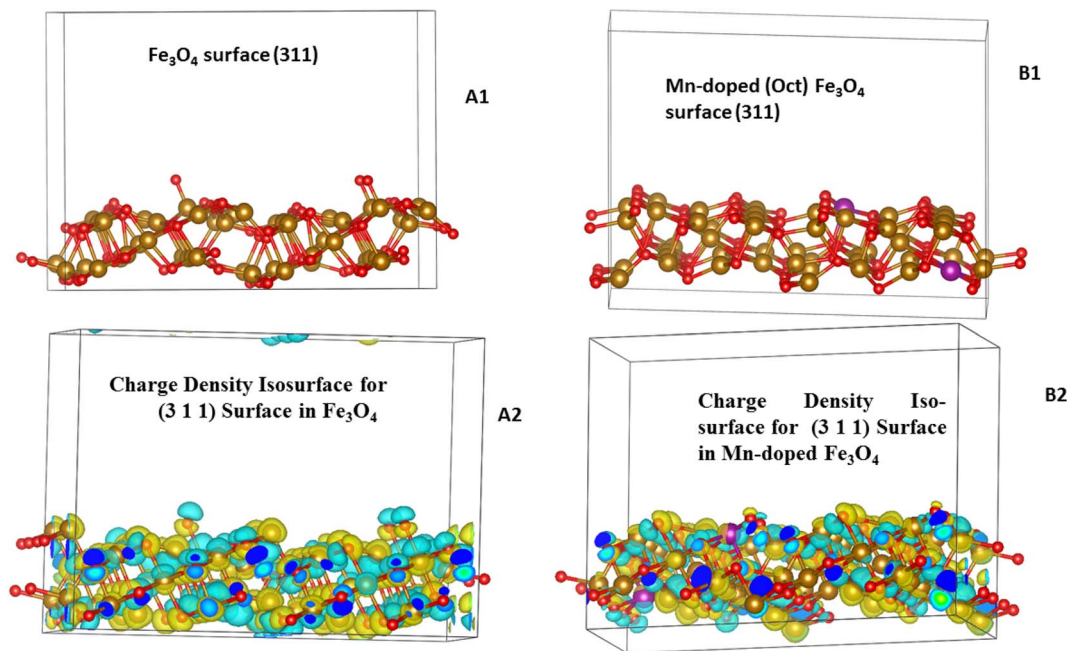


Fig. 16 (A1) Fe_3O_4 – (3 1 1) surface – 112 atoms with O in the top layer of the surface. (B1) Mn-doped Fe_3O_4 – (3 1 1) surface with 112 atoms with O in the top layer of the surface. Charge density iso-surface plot for the (3 1 1)-surface for (A2) Fe_3O_4 and (B2) Mn-doped Fe_3O_4 .

3.9.2 $[\text{UO}_2]^{2+}$ interaction with the (3 1 1) surface of Mn-doped Fe_3O_4 . The interactions with the uranyl ions (Fig. 17) were added to Mn, the Fe-octahedral site, and the Fe-tetrahedral site on the Mn-doped Fe_3O_4 (3 1 1) and were optimised for the energy calculations to determine the interaction energy as given in eqn (5).

$$E_{\text{Int.}} = E_{\chi-\text{UO}_2^{2+}} - E_{\text{slab}} - E_{\text{UO}_2^{2+}} \quad (5)$$

where $E_{\text{Int.}}$ = interaction energy, $E_{\chi-\text{UO}_2^{2+}}$ = the energy of $[\text{UO}_2]^{2+}$ ions adsorbed at a particular site (χ) in the Mn-doped Fe_3O_4 (3 1 1) surface, E_{slab} = slab energy of the Mn-doped Fe_3O_4 (3 1 1) surface, $E_{\text{UO}_2^{2+}}$ = energy of the $[\text{UO}_2]^{2+}$ ion. Note: S.I unit of Energy is J; (1 eV = 1.6021×10^{-19} J).

The interaction energy (eqn 5) as a measure of interactions between different sites of the Mn-doped Fe_3O_4 (3 1 1) is reflected in Table 3. The uranyl ion is shown in Fig. S4† and the energy for the optimisation of UO_2 and $[\text{UO}_2]^{2+}$ ion is shown in Table S6.†

Table 3 shows the interactions of the UO_2 -ion with different sites of the Mn-doped Fe_3O_4 (3 1 1) surface. The interaction with Mn (mostly Mn^{2+}) as indicated earlier, and also from the XPS studies in the experimental section, is the least. The results, therefore, corroborate the previous XPS results, which are shown in Scheme 6. However, it is worth mentioning that the surface as calculated with the *ab initio* technique is the surface with a vacuum above it and not ending in –H or –OH as will happen in the solvated state. However, in the solvated state, if it is considered that the solvation energy will be generically constant for the different sites for a particular surface and a particular ion, then it will be mainly governed by the interactions between the UO_2 -ions and the particular sites present over the surface as calculated herein. Therefore, both experimentally and theoretically, it is quite clear that the UO_2 -ions will be mainly adsorbed over the Mn^{2+} sites in the Mn-doped Fe_3O_4 surface, which is governed mainly by two factors. (1) Surface energy: Mn-doped Fe_3O_4 (3 1 1) < Fe_3O_4 (3 1 1). (2)

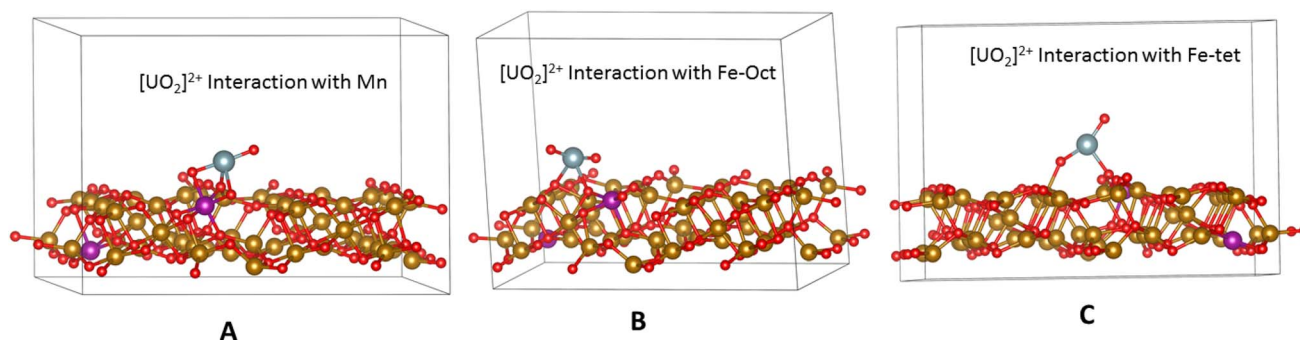


Fig. 17 Electronic interactions between $[\text{UO}_2]^{2+}$ and the (3 1 1) Mn-doped Fe_3O_4 surface, where the surface metal ion is present as (A) the Mn-octahedral site; the (3 1 1) Fe_3O_4 surface where the surface metal ion is present as (B) the Fe-octahedral site; (C) Fe-tetrahedral site.



Table 3 The interaction energies of uranyl ions with different sites in the Mn-doped Fe₃O₄ (311) surface (the complete calculation is given in the ESI)

Mn-doped Fe ₃ O ₄ -(311) surface sites	Mn-octahedral (eV)	Fe-octahedral (eV)	Fe-tetrahedral (eV)
[UO ₂] ²⁺ interaction energy	−17.048144	−16.29179	−16.269136

Interaction energy with UO₂-ion: Mn-interaction < Fe-interaction. As observed earlier, at the neutral pH (Fig. 12) the K_d value (Fig. S3†) initially increases from Fe₃O₄ to Mn-doped Fe₃O₄ and with higher concentrations of Mn-doping, it decreases after Mn-2. Both factors favour the Mn-doped Fe₃O₄ surface as compared to that of the Fe₃O₄ surface as indicated by experiment and *ab initio* calculations. However, for more Mn-doping, the interaction energy will play the dominant role in the surface energy. Regarding the interaction energy, the two factors (the slab energy of the Mn-doped Fe₃O₄ and the energy of the UO₂-ions) will be constant, and only the number of uranyl ion components will vary. However, the first term of eqn (4) will vary with different numbers of Mn-doped sites and will be the governing factor in explaining the $t_{1/2}$ or the K_d . Also, for different pH, the solvation has to be considered to explain this model completely.

The agglomeration/aggregation of charged particles and their interaction in liquid media is usually described by the DLVO (Derjaguin, Landau, Verwey, Overbeek) theory. This theory takes into account both van der Waals attraction and the electrostatic repulsion (double layer force).⁷⁹ Here, the electrostatic repulsion in the DLVO theory is due to the formation of a double layer. Here, the calculations were performed by taking into account the electrostatic interactions between Mn and different Fe sites over the Mn-doped Fe₃O₄ surface, considering that the solvation is the same for a particular surface and a particular ion at a given pH.

4. Conclusion

The sorption of uranyl ions over magnetic nanoparticles has garnered significant scientific attention due to its simplicity; however, clarification was needed regarding the exact sites of adsorption and the effective structural role played by the surface.

(a) This manuscript shows a simple, scalable route for synthesizing magnetic (Fe₃O₄) and doped magnetic nanoparticles (Mn-doped Fe₃O₄) for the sorption of uranyl ions. To the best of our understanding, these synthesized materials possess one of the best K_d ($\sim 3 \times 10^6$ cm³) values as compared to those reported in the literature for the different resins that are commercially used specifically for uranyl ions (*e.g.*, MnO₂ in polystyrene sulphonate resin $\sim 10^{6.1}$ or D-634 10^8 cm³);⁸⁰ however, the $t_{1/2}$ is perhaps the best with a value of 0.9 min. The stability of these MNP and Mn-MNPs in the entire pH range is also appreciable as compared to the literature.

(b) Mn²⁺ is substituted for the Fe²⁺ sites in the Fe₃O₄ lattice and as a result, the magnetic moment initially increases and then decreases. The alteration and the effect in sorption for the

textural properties are minimal and mainly result from the ionic interactions between the surface ions of the MNPs and the Mn-MNPs.

(c) In Fe₃O₄, the main interaction for the sorption of the uranyl ions is with the Fe²⁺ ions present in the Fe₃O₄ surface. Likewise, in Mn-MNPs, the results from the interactions of the uranyl ions with the substituted Mn²⁺ ions on the surface of Mn-MNPs are considerably lower as compared to the Fe sites in the Mn-MNPs.

(d) The surface energy and surface charge also play major roles in the alteration of the sorption properties of the nanoparticles. The ionic interactions between Mn²⁺, Fe²⁺ and Fe³⁺ mainly form the basis for ionic repulsion, which interacts with the solvated cation, thus directly affecting the sorption rate of the same. Taking the solvation parameter as a constant, it was proved that the surface energy of Mn-doped Fe₃O₄ was lower as compared to the Fe₃O₄ surface. Similarly, the interaction energy of uranyl ions with Mn²⁺ was lower as compared to Fe²⁺ and Fe³⁺ surface centres. The interplay of these two factors along with the surface charge is best optimised in neutral pH and plays different roles in the different pH ranges.

(e) This, in all probability, makes the Mn-2 sample the most optimised sample in the neutral pH regime. The acidic medium, though better as compared to the basic medium, is not as effective as the neutral medium.

The above understanding paves a new way for doping different ions into the Fe₃O₄ surface to further enhance the sorption efficacy. The present materials can be utilised for the removal of low-level uranyl ions from the different bioassays to develop a new methodology for the quantitative estimation of uranyl ions in urine bioassays.

Author's contribution

Prakash Mandal has performed the synthesis and characterisation for the MNPs and the Mn-MNPs and also has conducted all the sorption experiments and have plotted all the Figures and required Schematics. P. D. Sawant was consulted for the technical advice for sorption of uranyl ions in the bioassays and provided required support for conducting the experiments and giving suggestion during writing the manuscript. K. Bhattacharyya has conceptualised the present idea; completed the *ab initio* calculations and written the complete manuscript and has guided throughout this project.

Conflicts of interest

The authors declare no conflict of interest in the present manuscript.



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