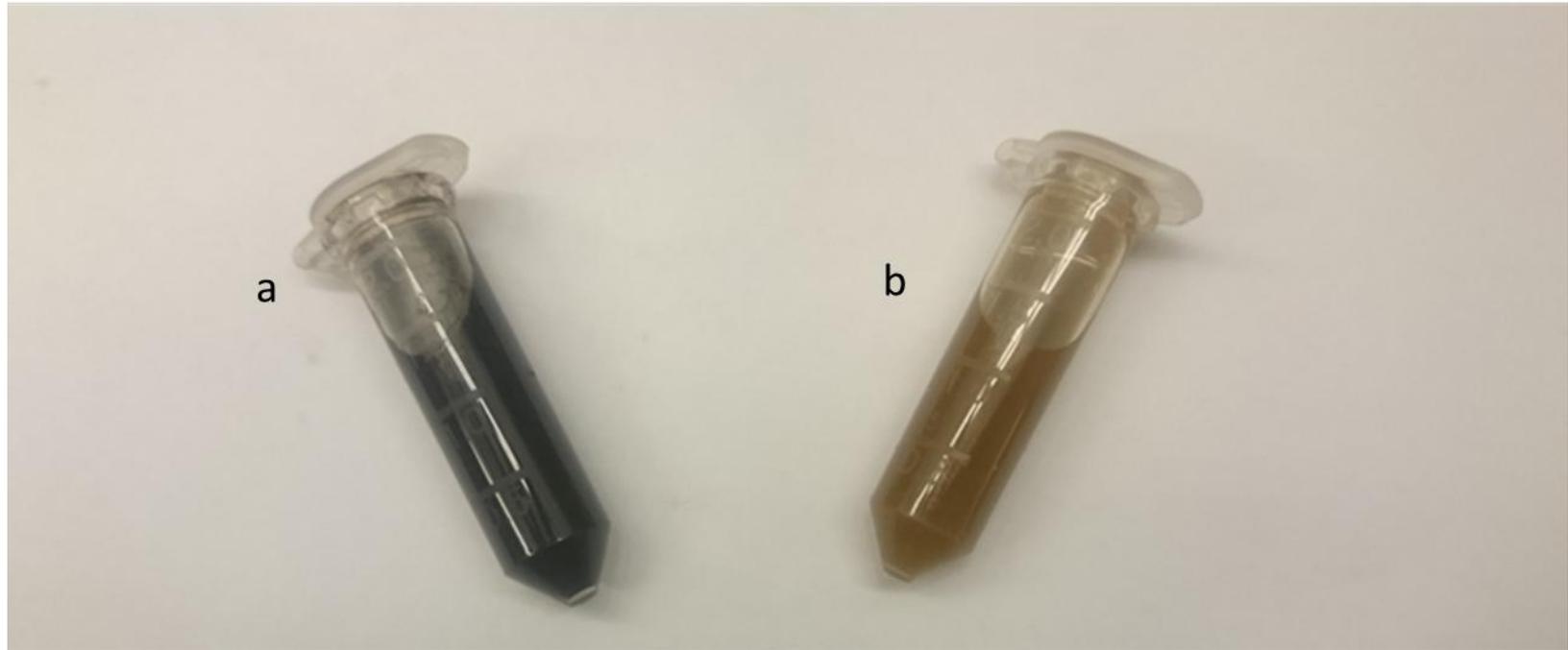


## Supplementary Information



**Supplementary information 1.** Change in color of MNPs as induced by changing their status from magnetite (a) to maghemite (b).

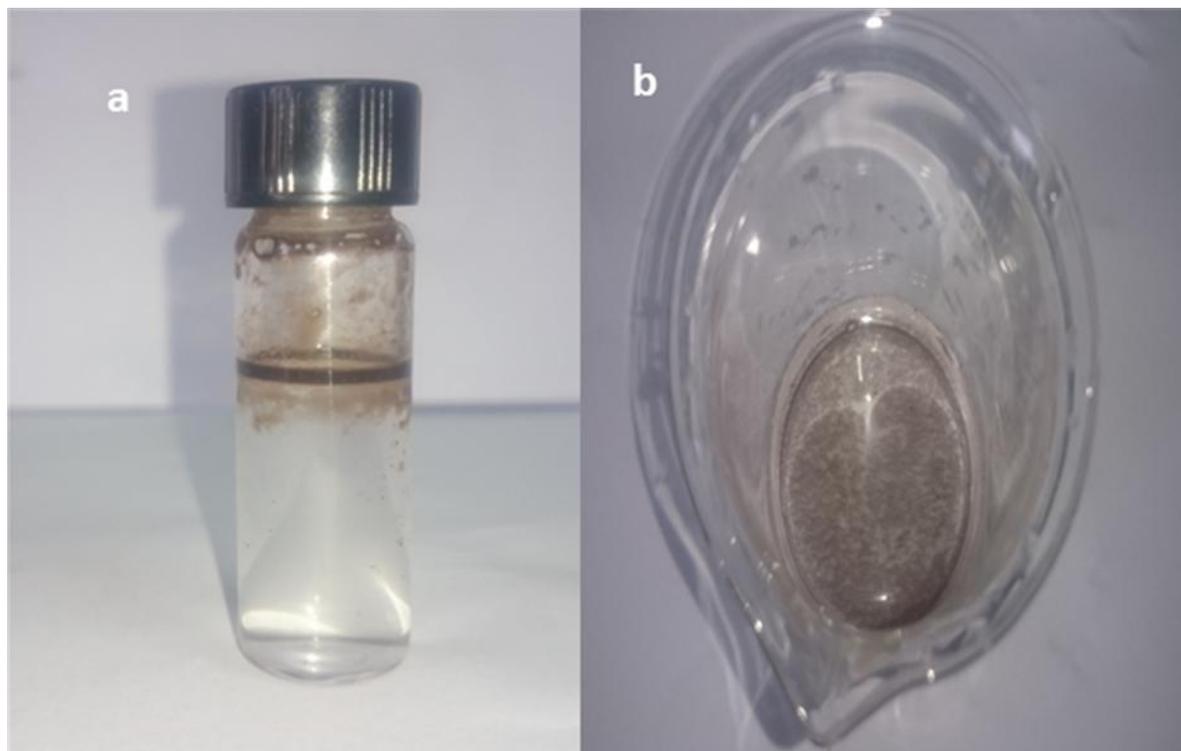
### Supplementary information 2. XRD data of the prepared MNPs

Position ( $2\theta$ )	$\text{\AA}(2\theta)$
30.29	2.957
35.72	2.518
43.38	2.091
57.45	1.605
63.07	1.474

\*D value: inter-atomic spacing in Angstroms ( $\text{\AA}$ )

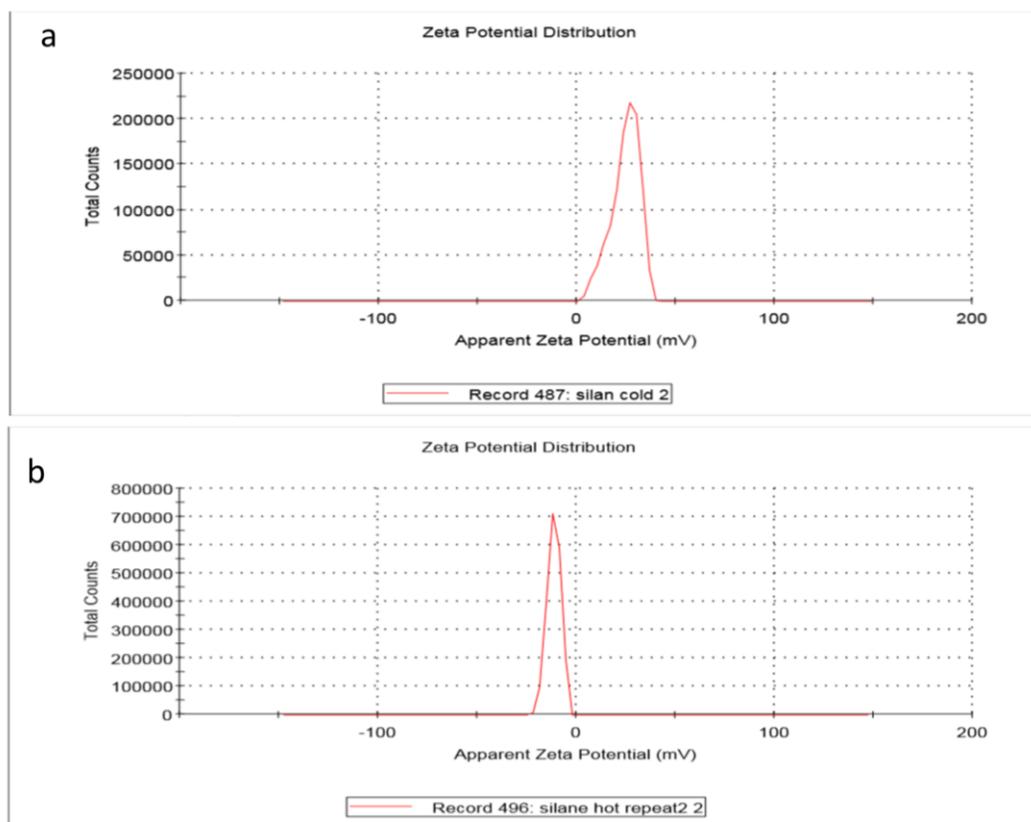
\*XRD was recorded at a Cu  $K\alpha$  anode ( $\lambda = 0.1542$  nm) operating at 40 kV and 30 mA. The pattern was collected at 25 °C in range 5 – 70° with a step size of 0.03° and step time of 1 s.

**Supplementary information 3.**



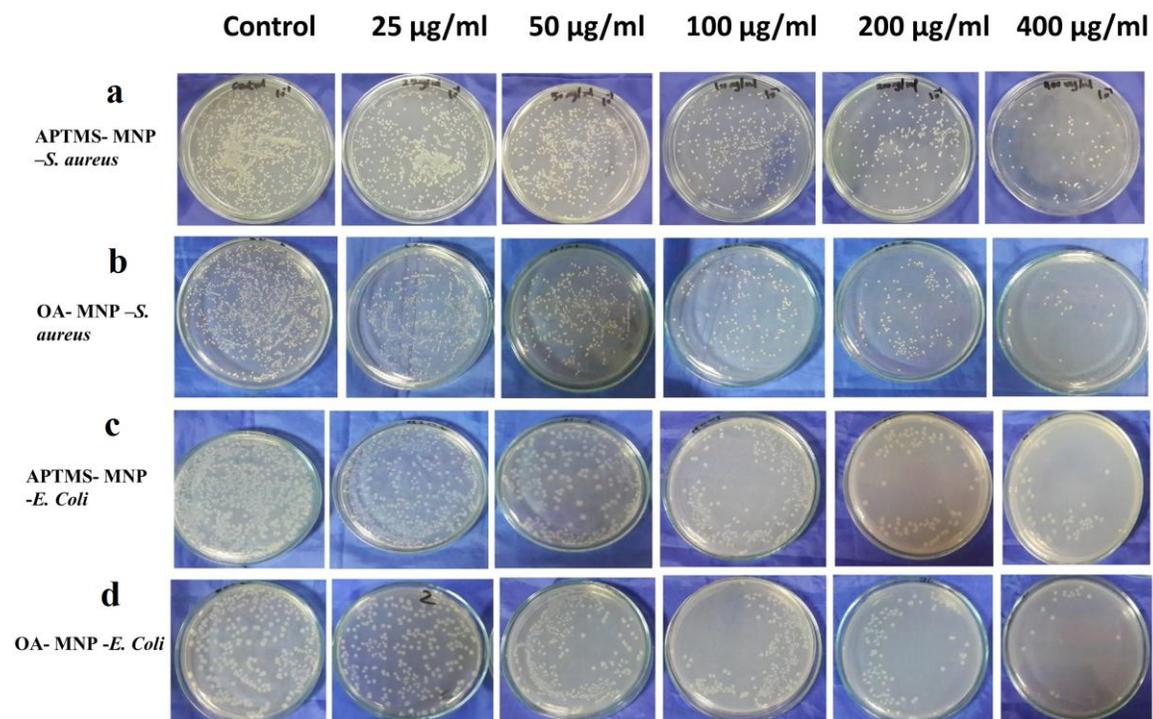
Supplementary information 3. (a) OA-MNP (after washing with water and before wash with ethanol) showing OA-NP not dispersed in water instead floating on water surface. (b) After successive washing with ethanol to remove excess OA from surface of MNPs; OA-MNP are still floating and unable to disperse.

#### Supplementary information 4.



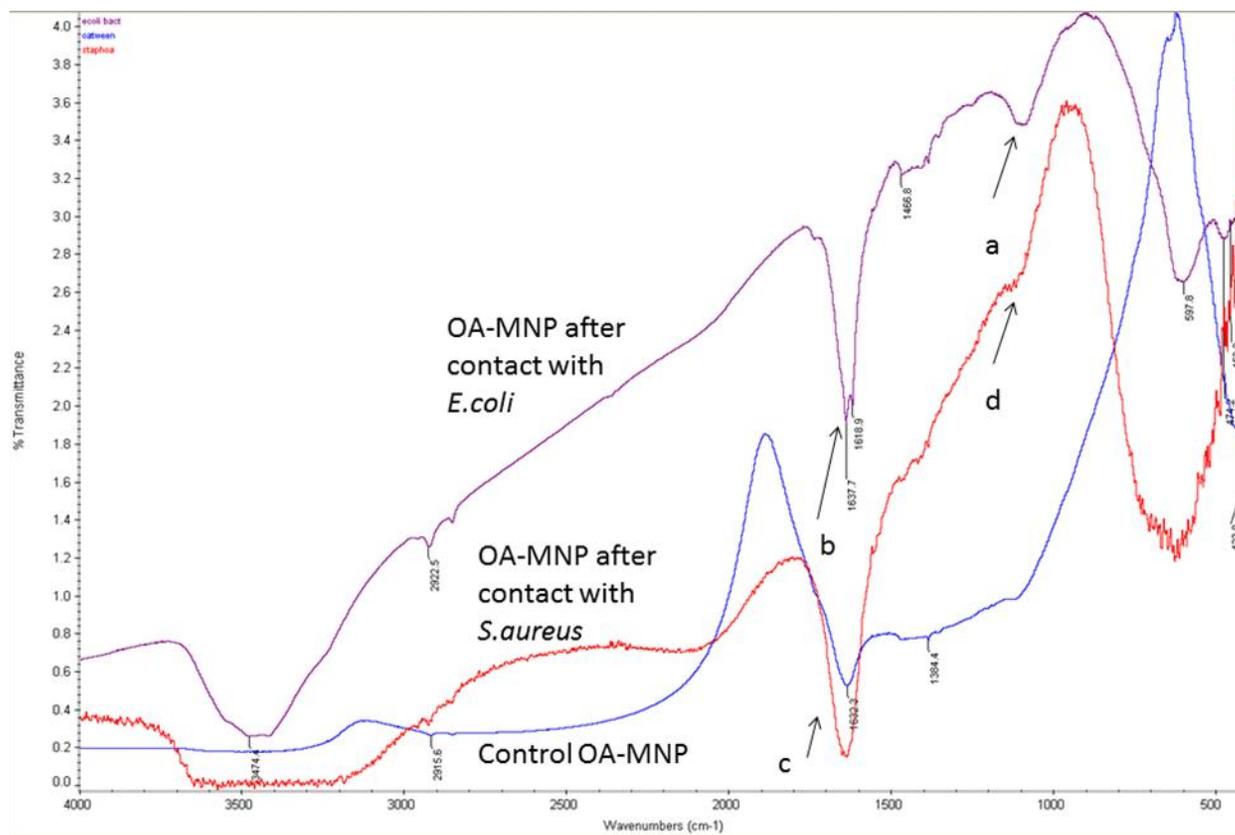
Supplementary information 4. Evaluation of the APTMS- MNP method was performed by comparing the Zeta potential shift imparted on the MNP. (a) Charge on surface of MNPs by the cold synthesis method (+25 mV). (b) Surface charge of particles prepared by the hot synthesis method (-11 mV). The cold synthesis method was more efficient to coat the MNPs with a surface positive charge.

## Supplementary information 5



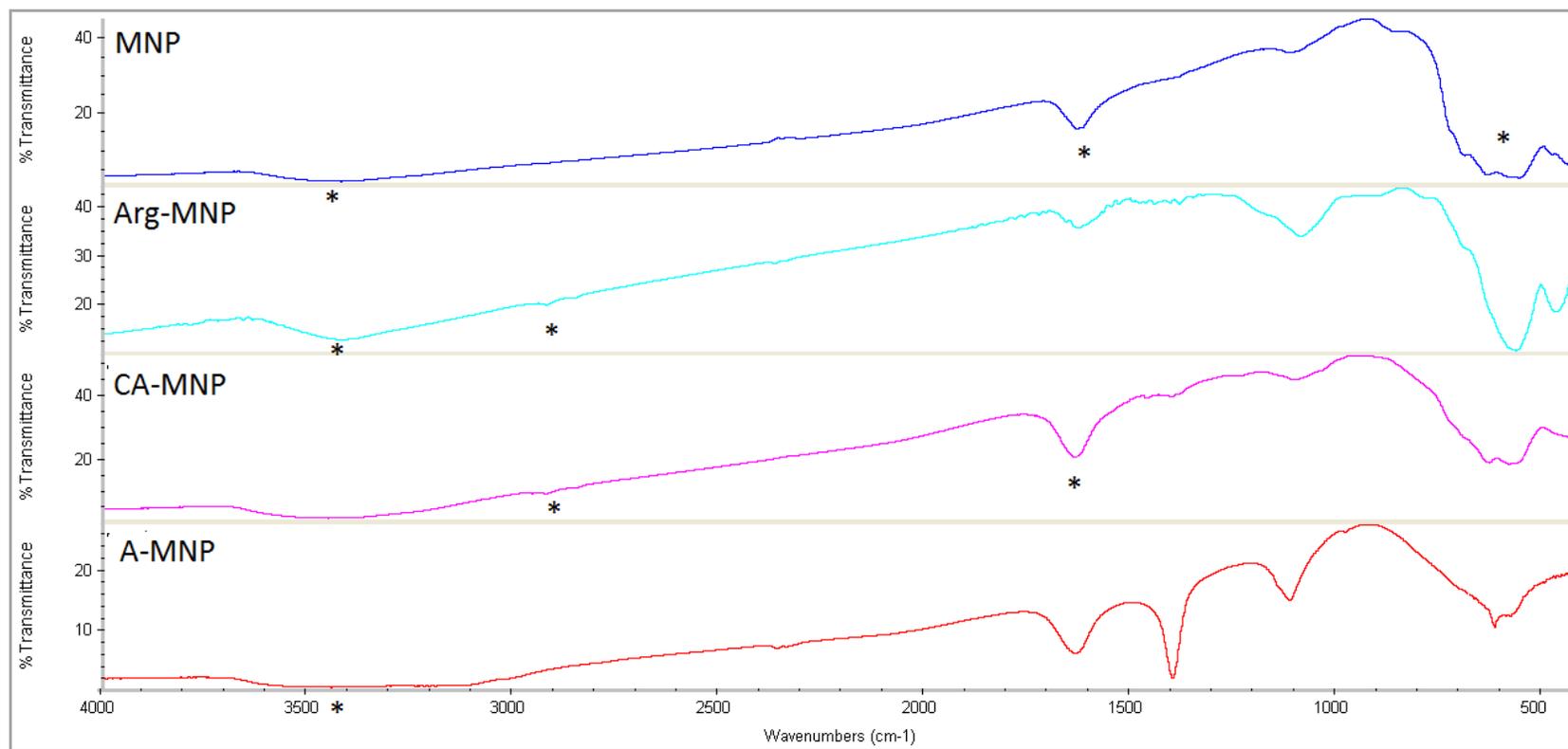
Supplementary information 5. Growth Inhibitory effect of MNPs on test bacteria 24 h post incubation with variable concentrations of MNPs. Subculture *S.aureus* (ATCC-6538) and *E.coli* (ATCC-8739) were incubated with varying concentrations of APTMS-MNP and OA-MNP for 24 h at 37°C on a shaker (200 rpm) for 24 h. At the end of the incubation period, samples were obtained from each flask and 10 fold serially diluted in sterile saline. One hundred  $\mu\text{L}$  of each dilution as well as control were spread on the surface of 3 nutrient agar plates, incubated at 37°C for 24 h and the average numbers of colony forming units (CFU/mL) were counted. (a) and (c) show the effect of APTMS-MNP on *S. aureus* and *E. coli*. (b) and (d) show the effect of OA-MNP on *S. aureus* and *E. coli*.

## Supplementary information 6.



Supplementary information 6. FTIR spectrum of control OA-MNPs before and after contact with *S. aureus* and *E. coli*. For *E. coli* (a) indicates the polysaccharide (900 -1200 cm<sup>-1</sup>) and (b) indicates the band attributed to primary amine (1640 – 1560 cm<sup>-1</sup>) [31]. For *S. aureus*, the peaks denoted by (c) and (d) represent the C=N and the C-H, respectively [32, 33].

### Supplementary information 7.



**Supplementary information 7. FTIR spectra of MNPs which did not show significant antibacterial potentials. (\*) represent characterizing band.**