

Supplementary Information

Size-dependent antimycobacterial activity of titanium oxide nanoparticles against

Mycobacterium tuberculosis

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Table S1 UV and DRS UV, binding energy, Zeta potential and size distribution, average size (TEM and XRD)

TiO₂	Bandgap (eV)	Zeta potential (mv)	Polydispersity index (PDI)	Size distribution (nm)	Average size (nm)	
					TEM	XRD
6 hr	3.9	-12	0.712	70-85	77.4	75.8
12 hr	3.56	-24	0.284	45-60	53.5	53.2
18 hr	3.21	-30	0.139	26-38	31	30.4
24 hr	3.06	-38	0.057	10-22	16	16.5

Fig. S1 UV-visible diffuse reflectance (DRS) spectra of TiO₂ nanoparticles (a) and the Tauc plot (b)

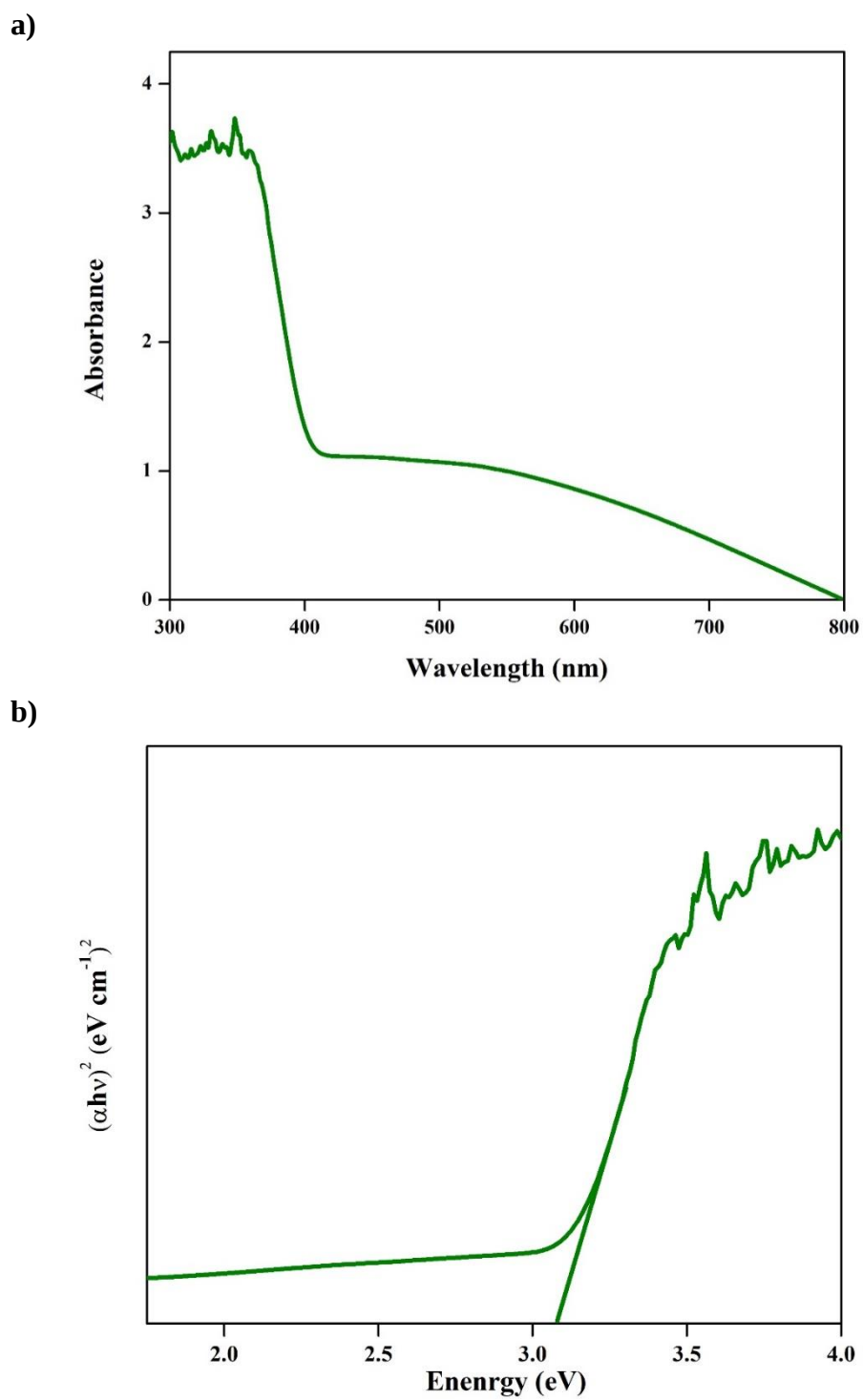
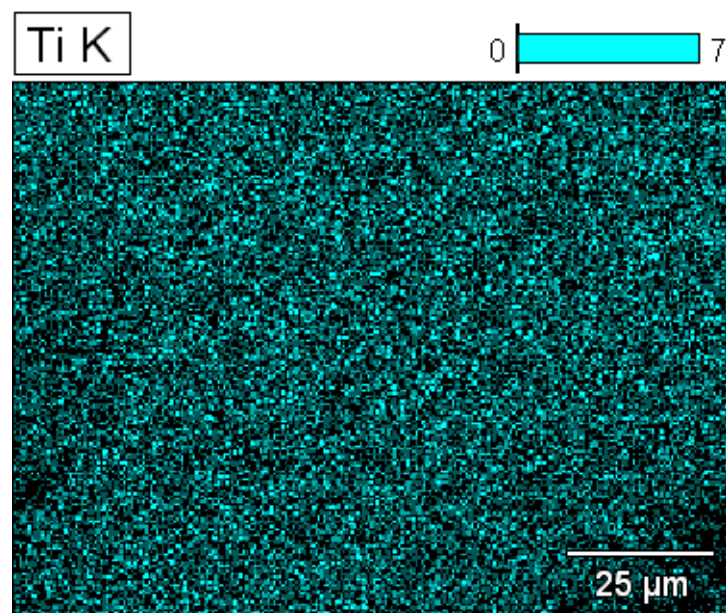


Fig. S2 the elemental mapping of TiO_2 nanoparticles observed using SAM nano probe for Ti (a), and O (b).

a)



b)

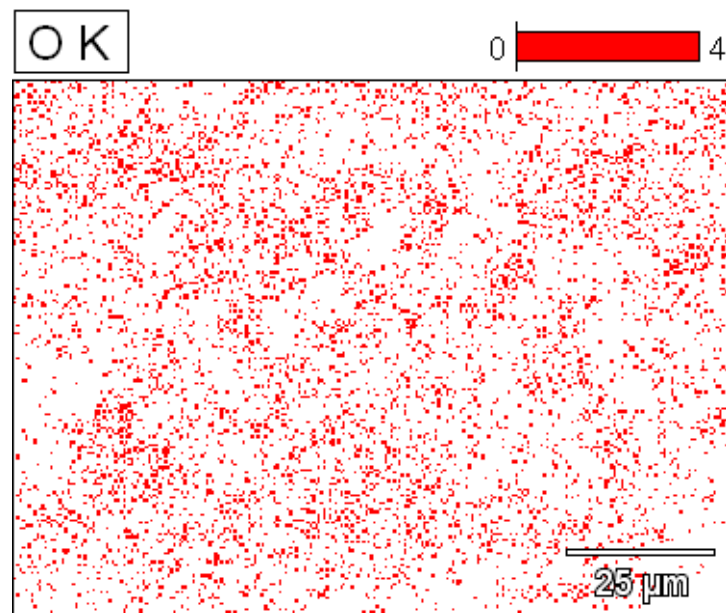
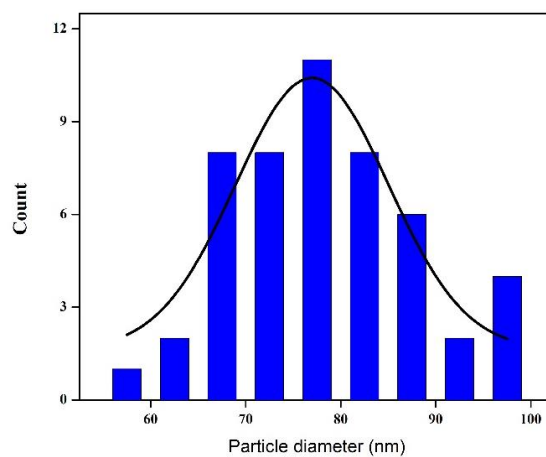
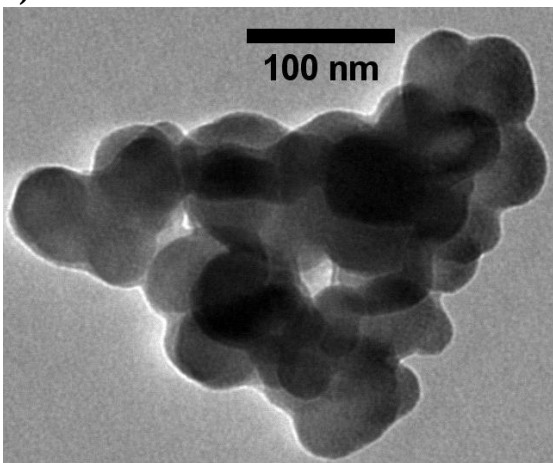
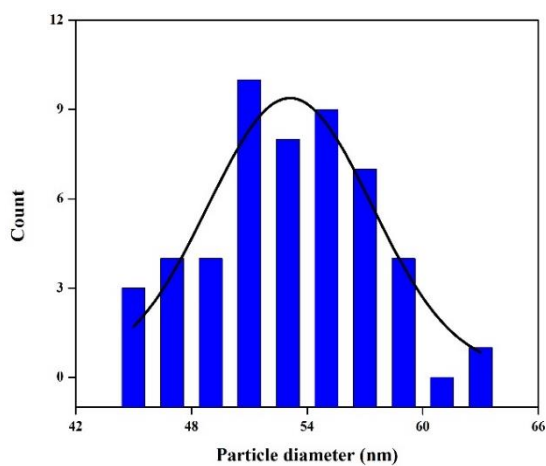
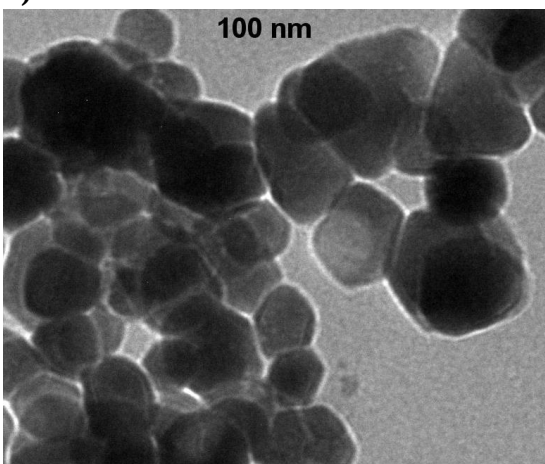


Fig. S3 TEM micrographs of TiO_2 and its corresponding size distribution histogram prepared at different time interval 6h, 12h and 18h

a)



b)



c)

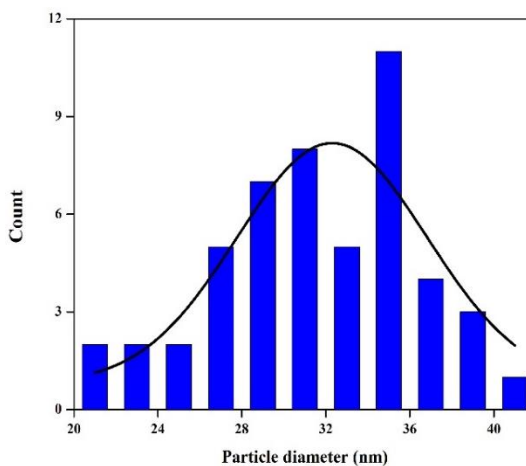
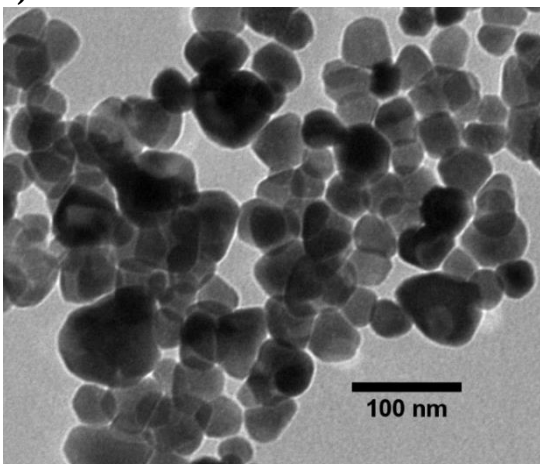


Fig. S4 Zeta potential of TiO₂ nanoparticles prepared at different time interval 6 h (70-85 nm), 12 h (45-60 nm), 18 h (26-38 nm) and 24 h (10-22 nm)

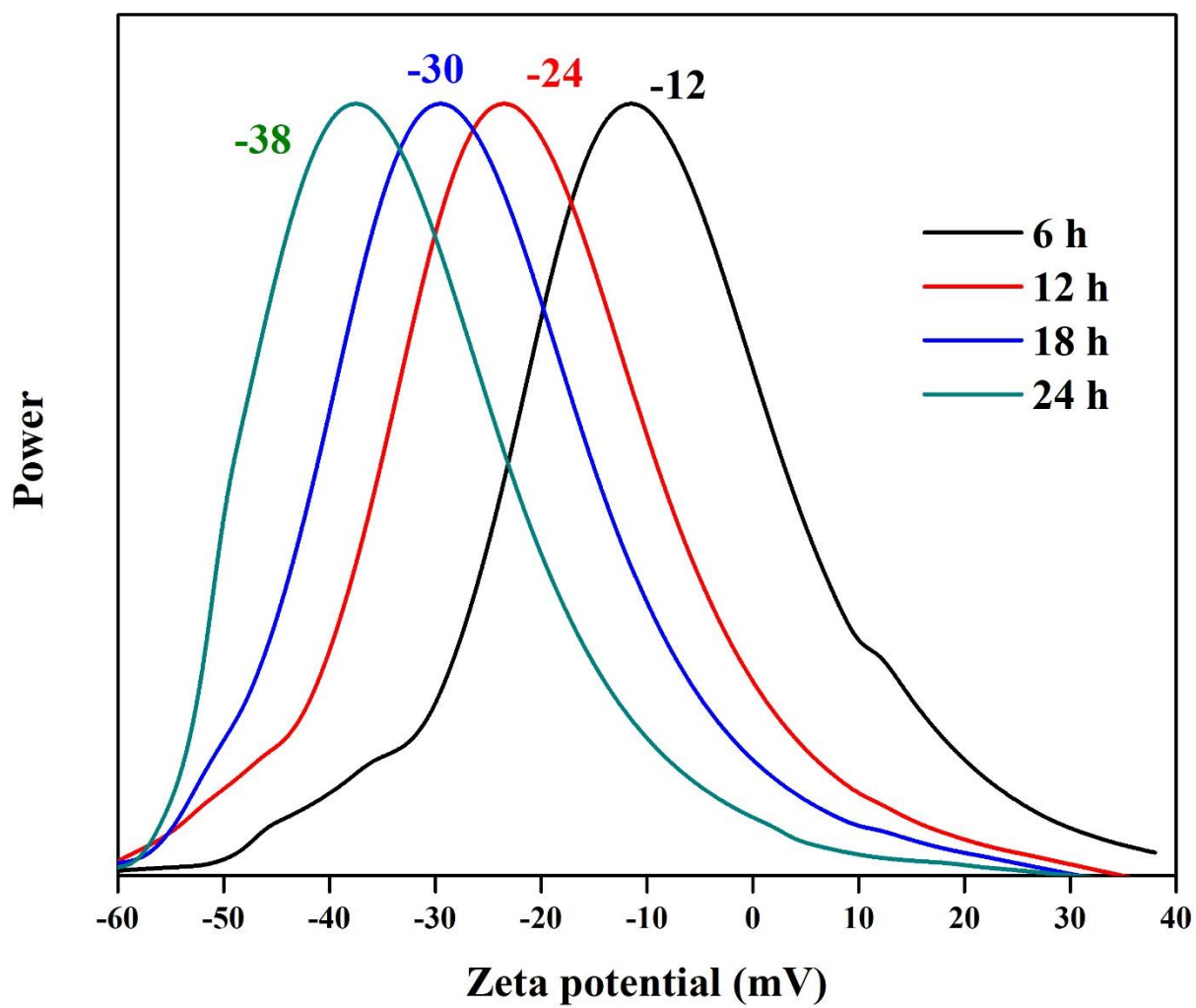


Fig. S5 The antimycobacterial activity of Rifampicin (0-20 $\mu\text{g/ml}$) against *M. tuberculosis* (a), *M. bovis* (b) and *Mycobacterium* sp. (c) after 24 h treatment.

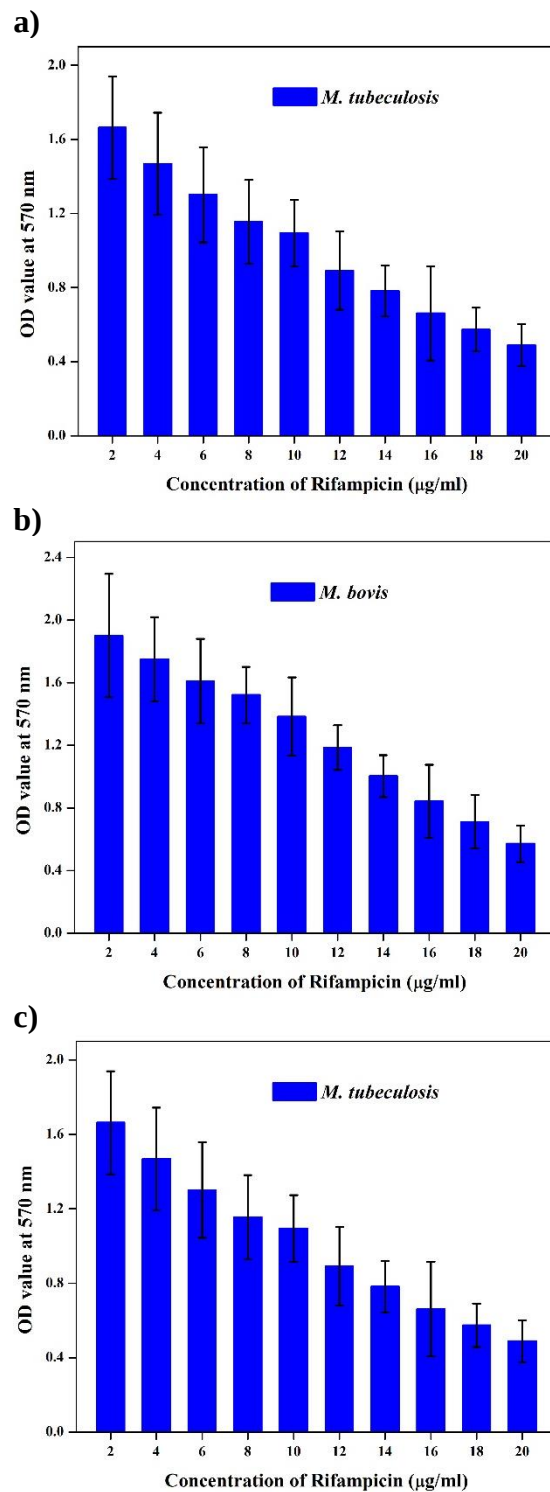


Fig. S6 Effect of 70-85 nm in size ranged TiO₂ nanoparticles (100 µg/ml) on biofilm formation of *M. tuberculosis*, *M. bovis* and *Mycobacterium* sp.

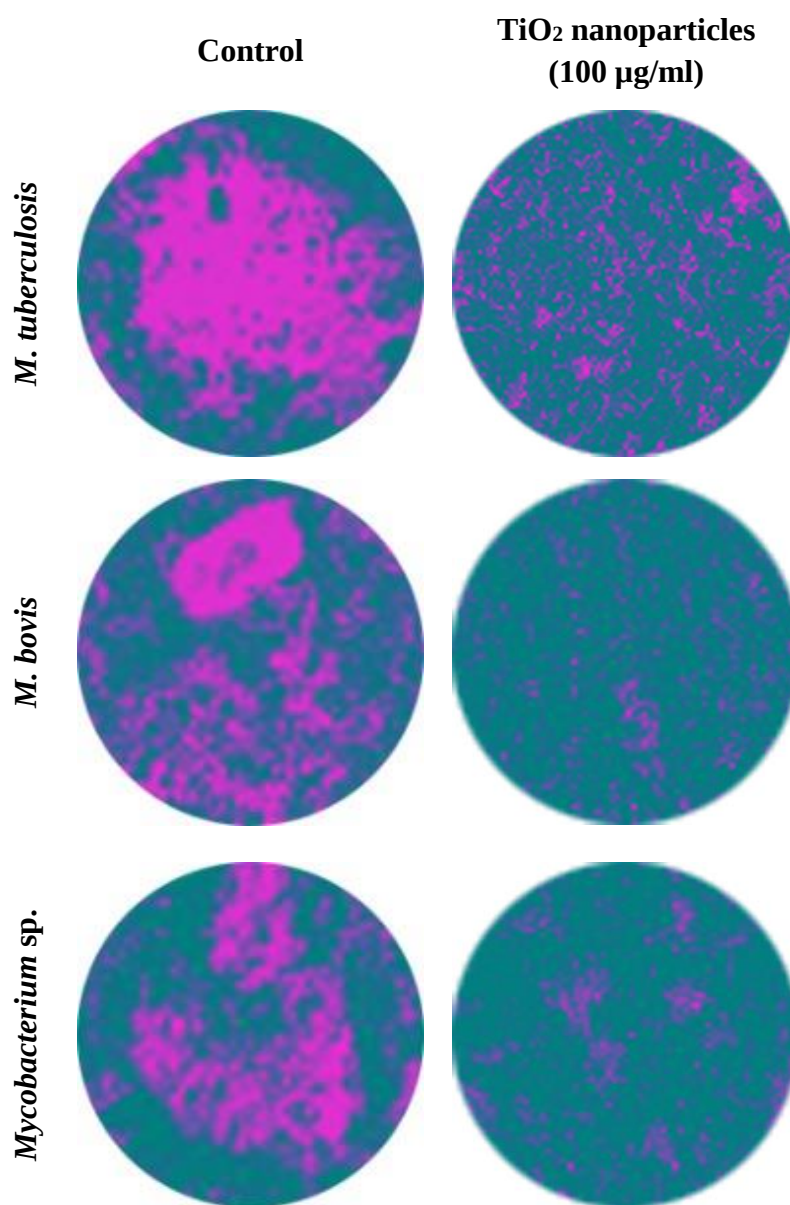


Fig. S7 Effect of 45-60 nm in size ranged TiO₂ nanoparticles (100 µg/ml) on biofilm formation of *M. tuberculosis*, *M. bovis* and *Mycobacterium* sp.

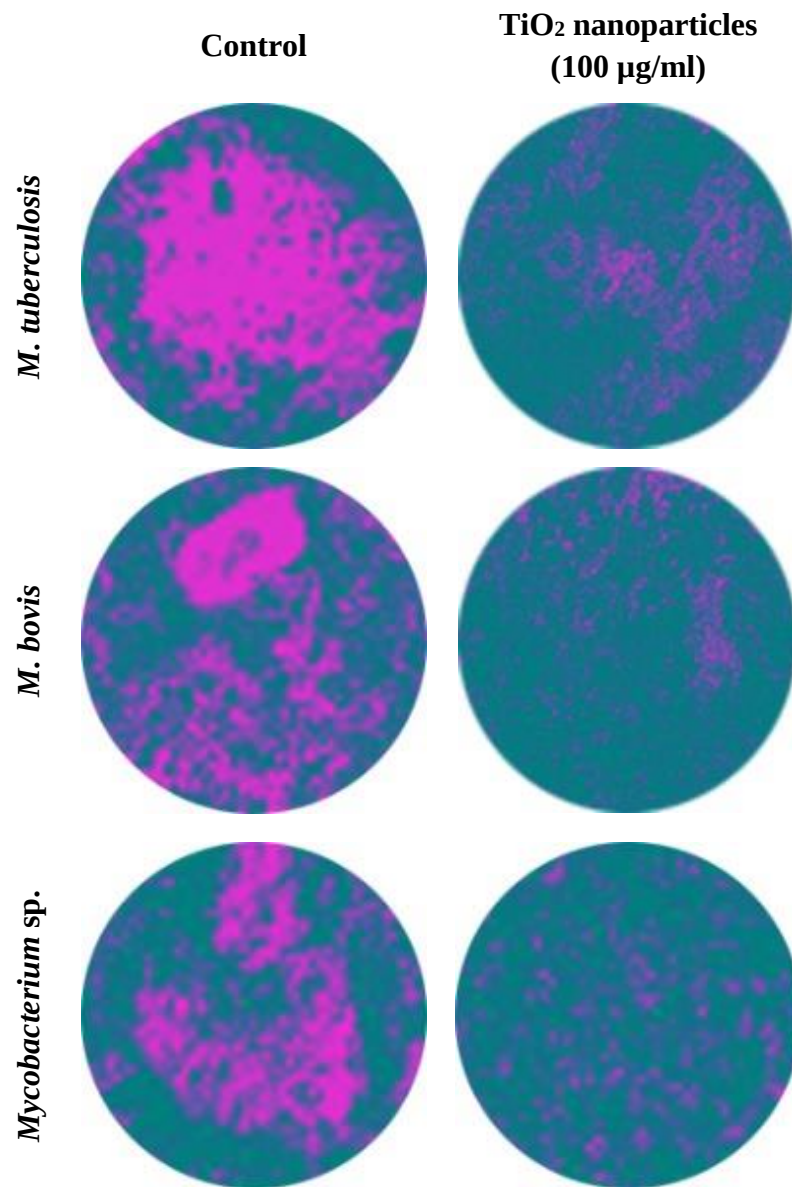


Fig. S8 Effect of 26-38 nm in size ranged TiO₂ nanoparticles (100 µg/ml) on biofilm formation of *M. tuberculosis*, *M. bovis* and *Mycobacterium* sp.

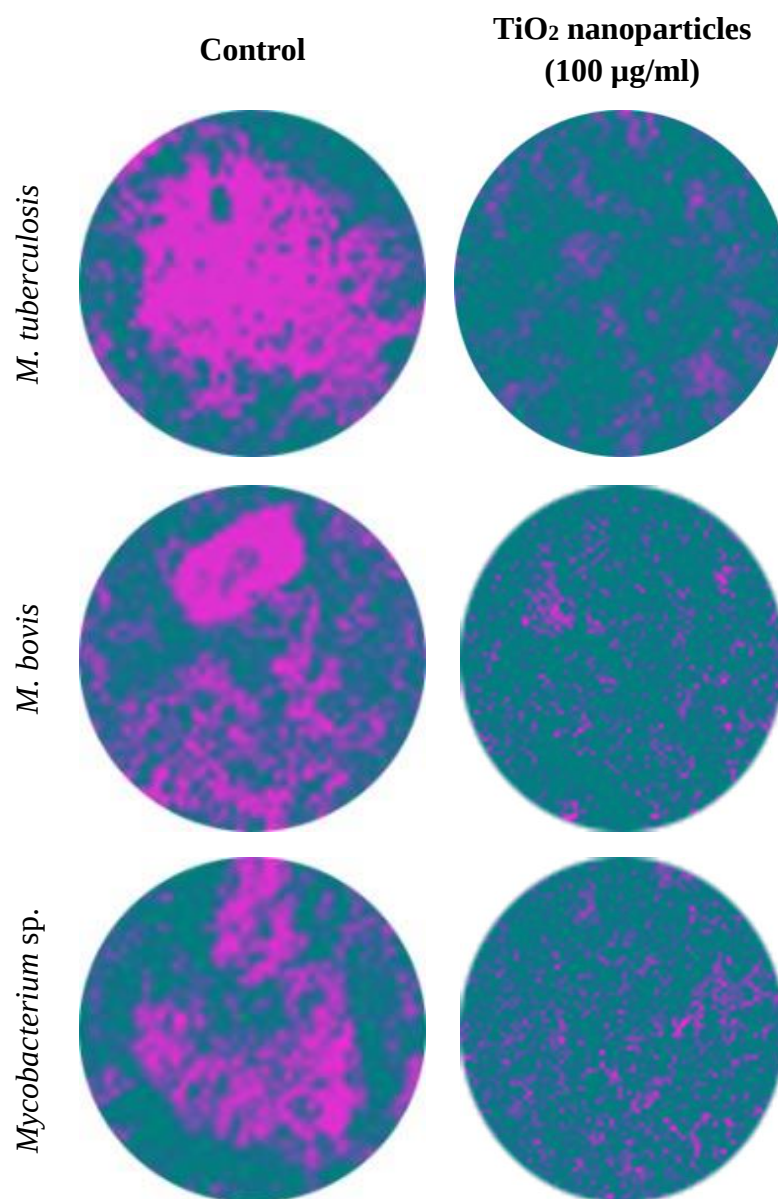


Fig. S9 Effect of Rifampicin (12 $\mu\text{g/ml}$) on biofilm formation of *M. tuberculosis*, *M. bovis* and *Mycobacterium* sp.

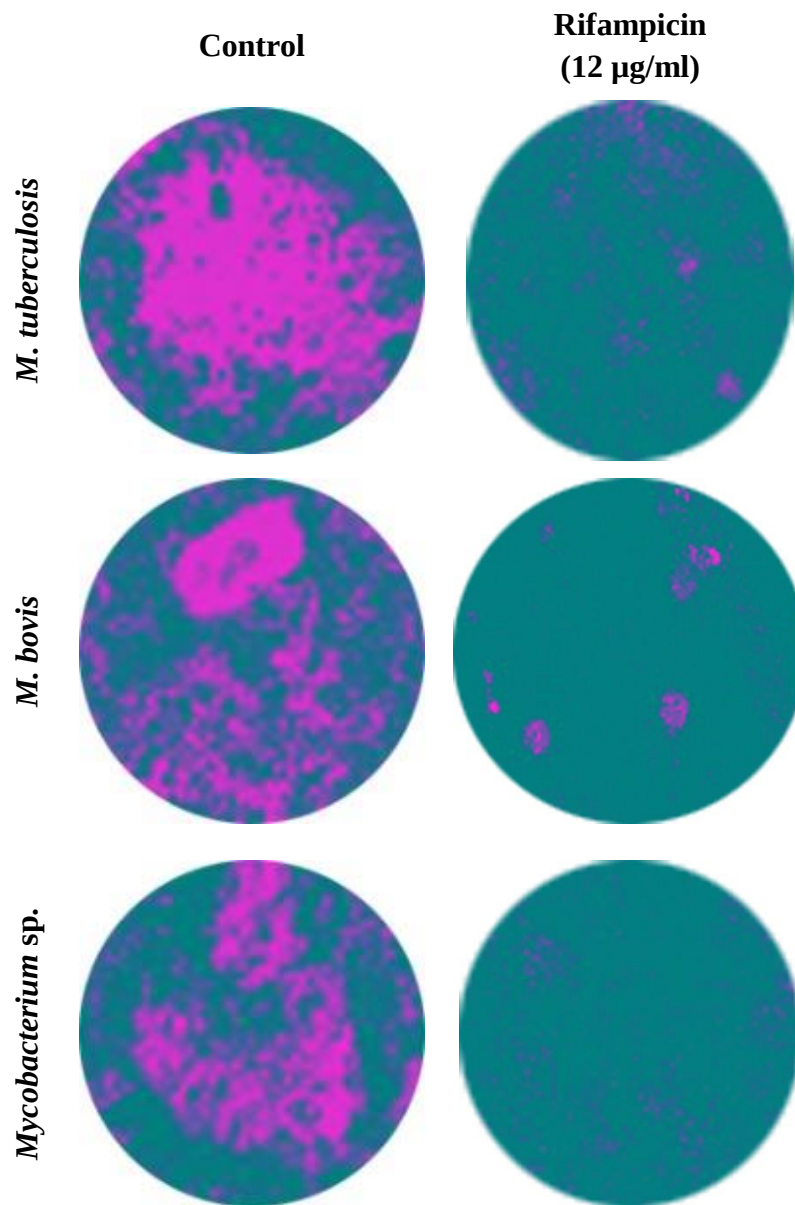


Fig. S10 Possible mechanism involved in the TiO_2 nanoparticles mediated inhibition of growth of mycobacteria

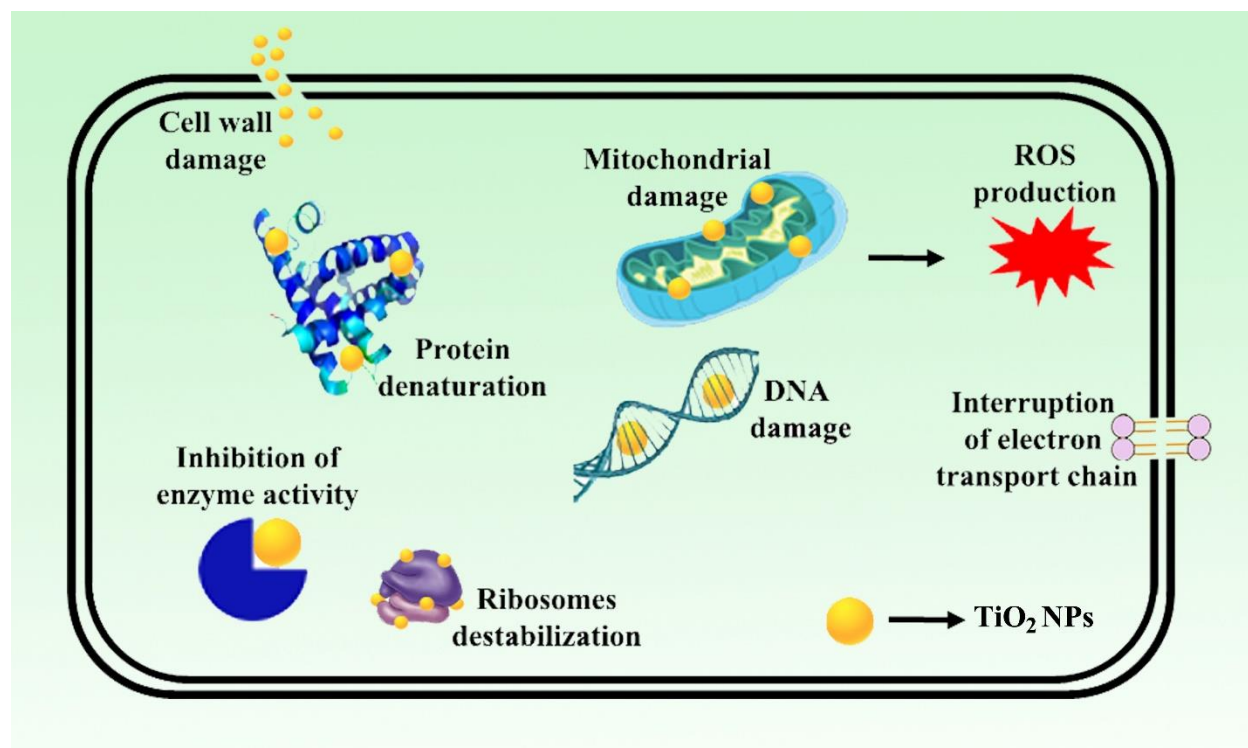


Fig. S11 *In vitro* toxicity of TiO₂ nanoparticles against normal human lung cells BEAS-2B (a) and the morphological changes in control (b) and TiO₂ nanoparticles treated cells

