

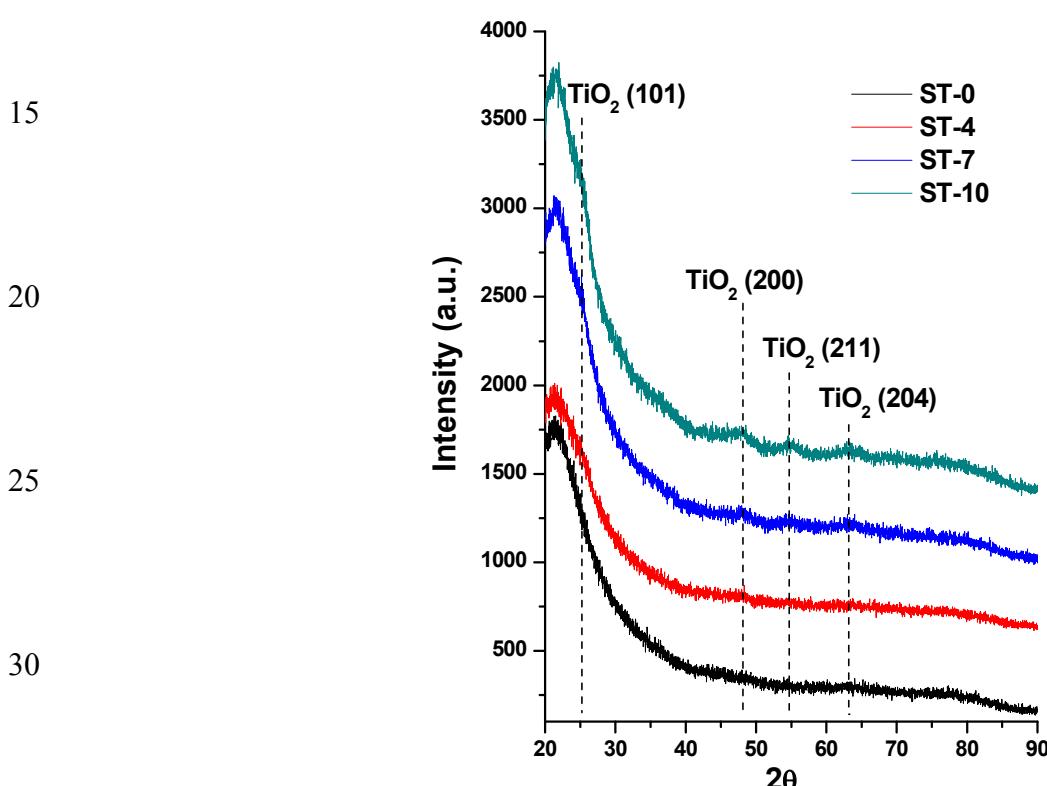
TiO₂ nanoparticles doped SiO₂ films with ordered mesopore channels: A catalytic nanoreactor†

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Electronic Supplementary Information (ESI)

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Fig. S1 High angle XRD patterns of TiO₂ doped SiO₂ films heat-treated at 350 °C. A grazing incidence angle of 0.3° was used to record the XRD patterns.

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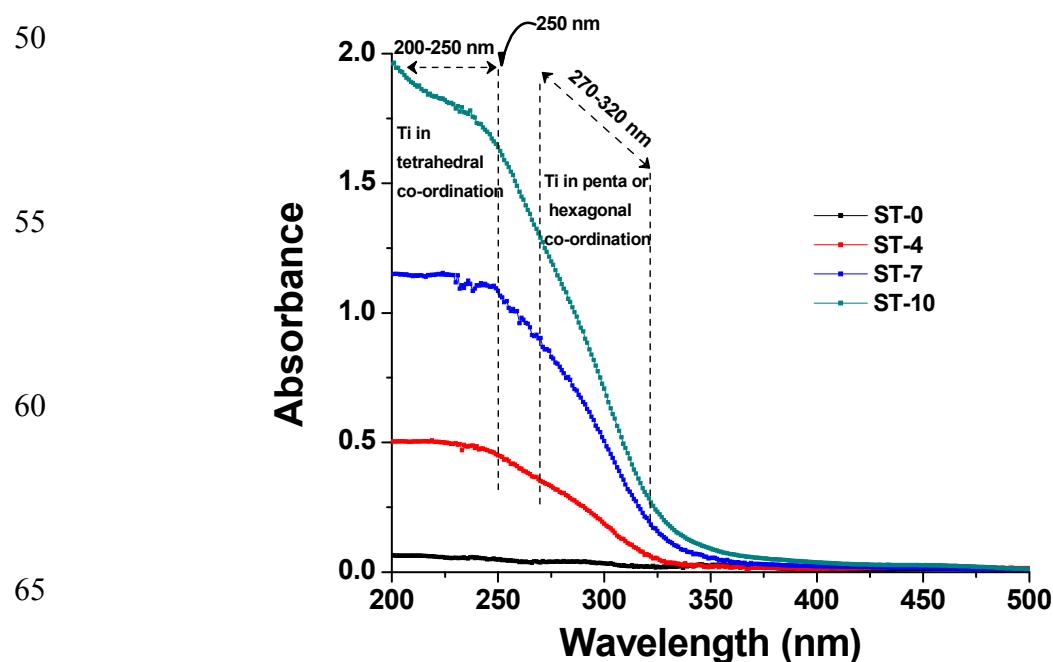
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70 **Fig. S2** UV-visible absorption spectra of TiO_2 doped SiO_2 films after heat-treatment at 350 °C; the peak assignments are given in the body of the figure.

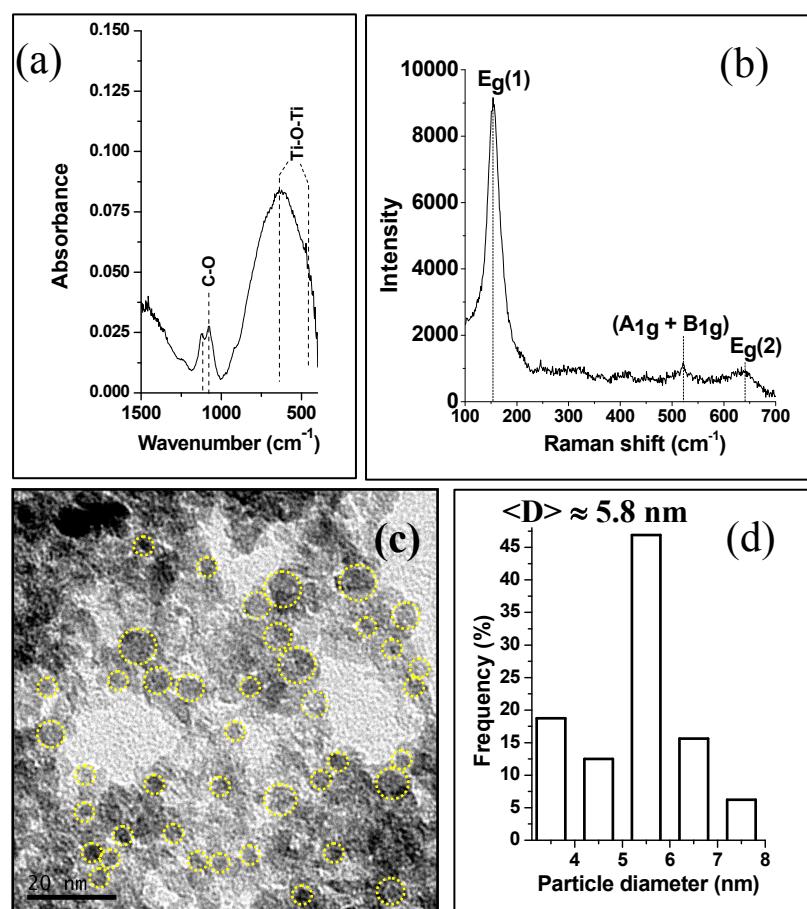
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Fig. S3 (a) FTIR, (b) Raman and (c) TEM images of TiO₂ NPs dispersion used in this work. The particle size distribution (average size $\langle D \rangle \approx 5.8 \text{ nm}$) evaluated from TEM is shown in (d).

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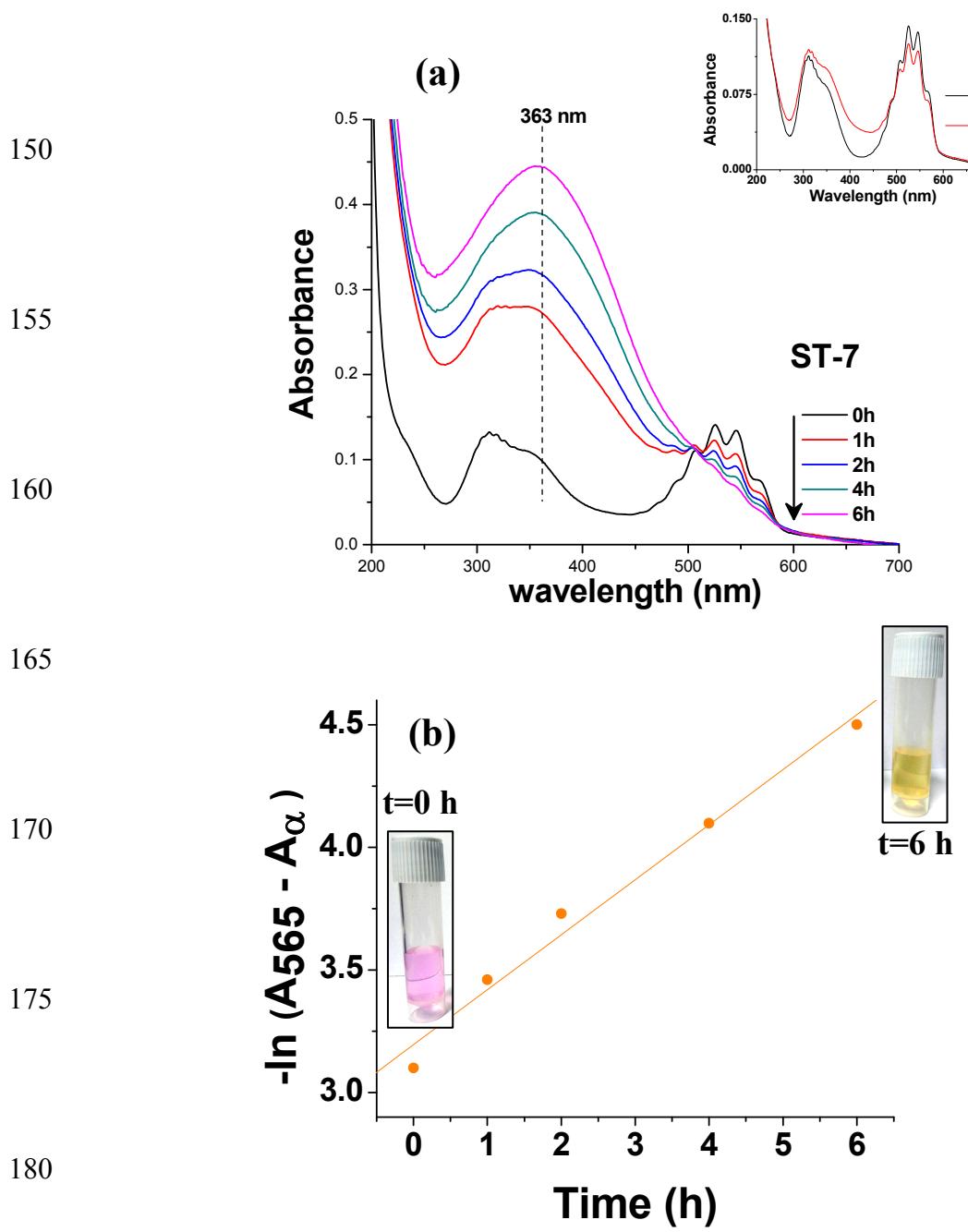


Fig. S4 (a) Evolution of UV-visible absorption spectra during the reduction of KMnO_4 in presence of 185 ST-7 (7 mol% TiO_2 doped SiO_2) film catalyst at 25°C ; inset shows the UV-visible spectra of the reaction in absence of film catalyst; (b) pseudo first order plot of $-\ln(A_{565} - A_\alpha)$ versus time for the reaction and inset shows the photos of initial (KMnO_4) and final (MnO_2) solutions.

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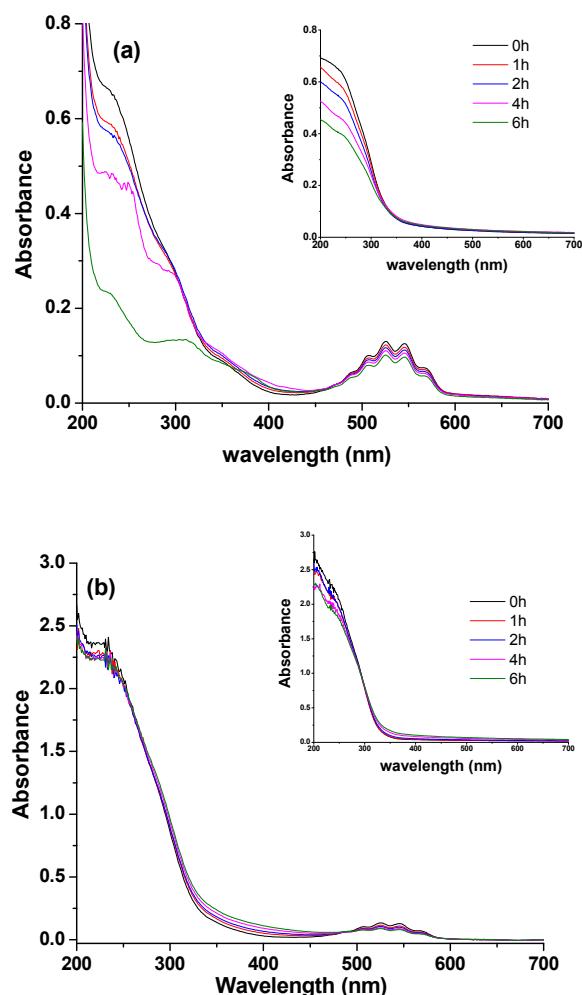


Fig. S5 UV-visible spectral evolution of aqueous KMnO_4 solution in presence of TiO_2 dispersion used 225 in this work: (a) equivalent amount of TiO_2 present in the ST-7 films used for catalytic reaction and (b) 4 times concentrated TiO_2 with respect to (a). The respective insects show the spectral changes of TiO_2 dispersion in water with time in absence of KMnO_4 .