

**Supporting information for**

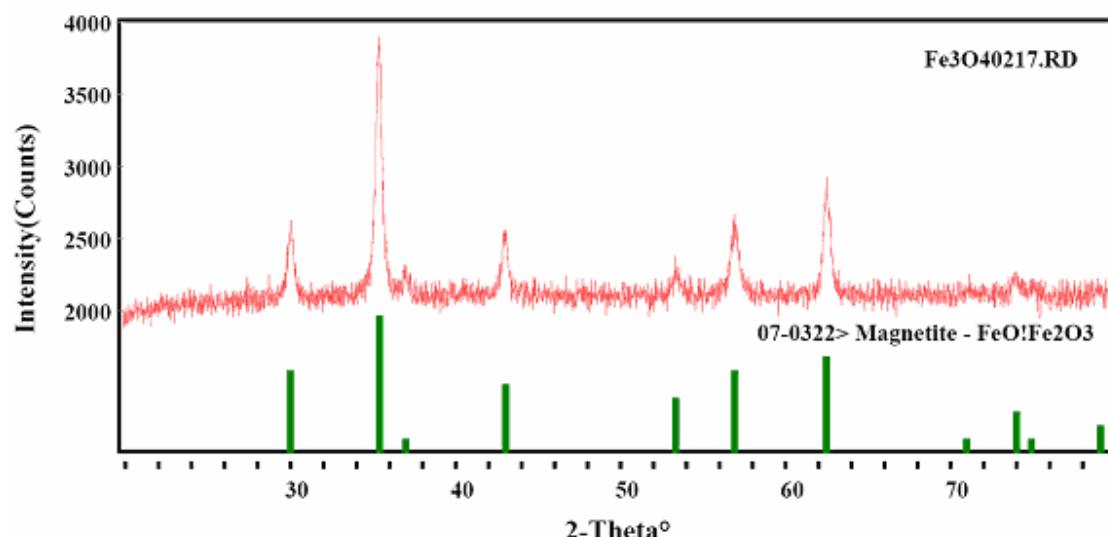
**Preparation of Hybrid Polymer Nanocomposite Microparticles  
by A Nanoparticle Stabilised Dispersion Polymerisation**

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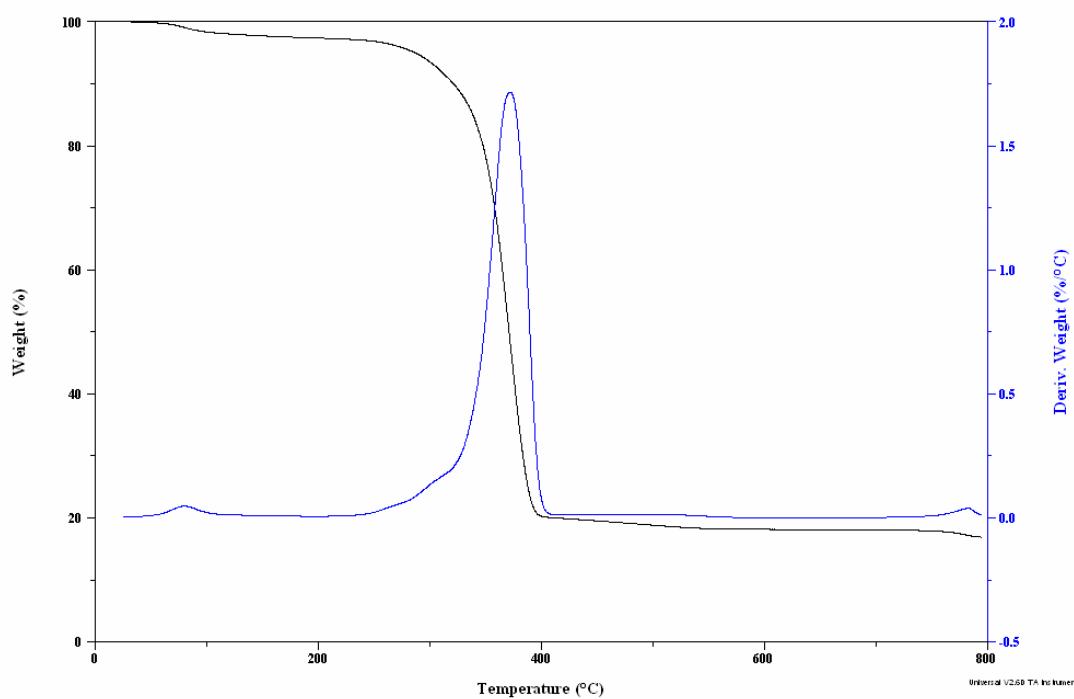
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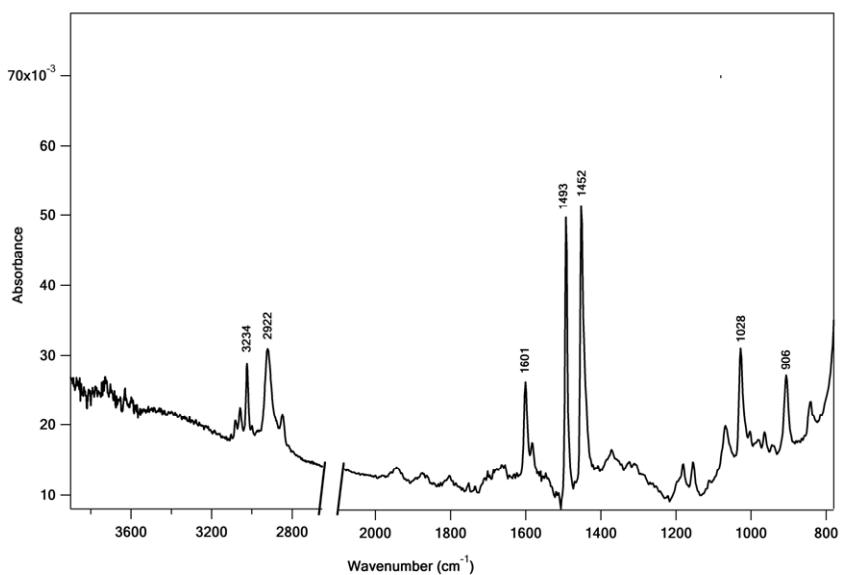


**Fig. S1 X-ray diffraction (XRD) spectrum of the Fe<sub>3</sub>O<sub>4</sub> nanoparticles derived from their aqueous solution which was used in this work.**

A suspension containing nanoparticles was allowed to settle for 12 hours and approximately 80% of the excess water was decanted. The remaining solution was frozen by liquid nitrogen and freeze dried at a temperature of -50°C to yield powder. The powder was characterised by Phillips EXPERT 1830 diffractometer with Cu K $\alpha$  radiation ( $2\theta$  0.02 degree/s from 20-80°). The obtained diffraction data were compared with the Joint Committee on Powder Diffraction Standards (JCPDS) - International Centre for Diffraction Data (ICDD) [07-0322]. For the main peak, the XRD estimated size is 22 nm based on the Scherrer equation.



**Fig. S2** TGA data of the Fe<sub>3</sub>O<sub>4</sub>-polystyrene nanocomposite synthesized in this work. The measurement was taken under flowing nitrogen at a heating rate of 5°C/min. The data clearly show the nanocomposite decomposes at 300-400°C and the total Fe<sub>3</sub>O<sub>4</sub> uptake is ca. 20%.



**Fig. S3** ATR-FTIR spectrum of the  $\text{Fe}_3\text{O}_4$ -polystyrene nanocomposite powder synthesized in this work. The major characteristic bands are labelled on the spectrum. The existence of  $\text{Fe}_3\text{O}_4$  nanoparticles capped on polystyrene partly impairs its infrared signal. However, the main peaks of the nanocomposite are not at all shifted from those of pure polystyrene, supporting strongly that there is no chemical interaction between  $\text{Fe}_3\text{O}_4$  and polystyrene.