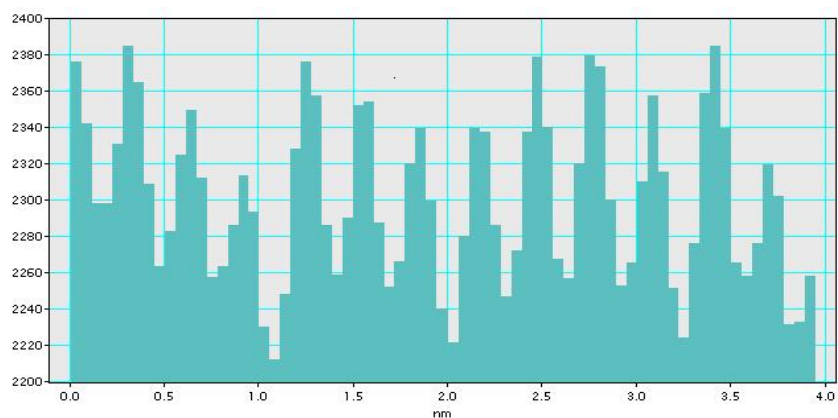
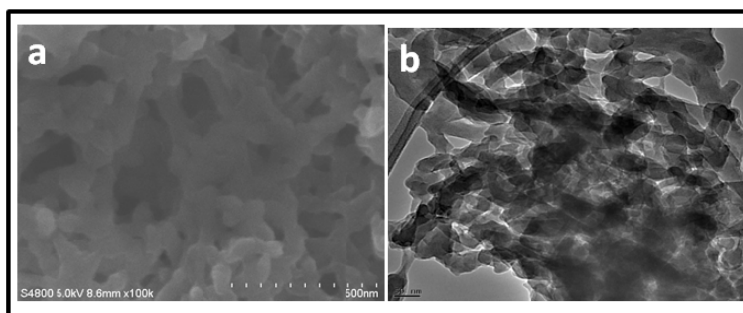


SI₁ – d spacing of PANI anchored SVO NC synthesized at 80⁰C

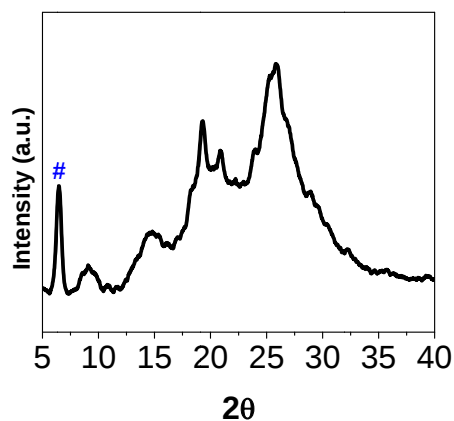


SI₂ a) FESEM b) TEM images of PANI-NF's synthesized at 60⁰C



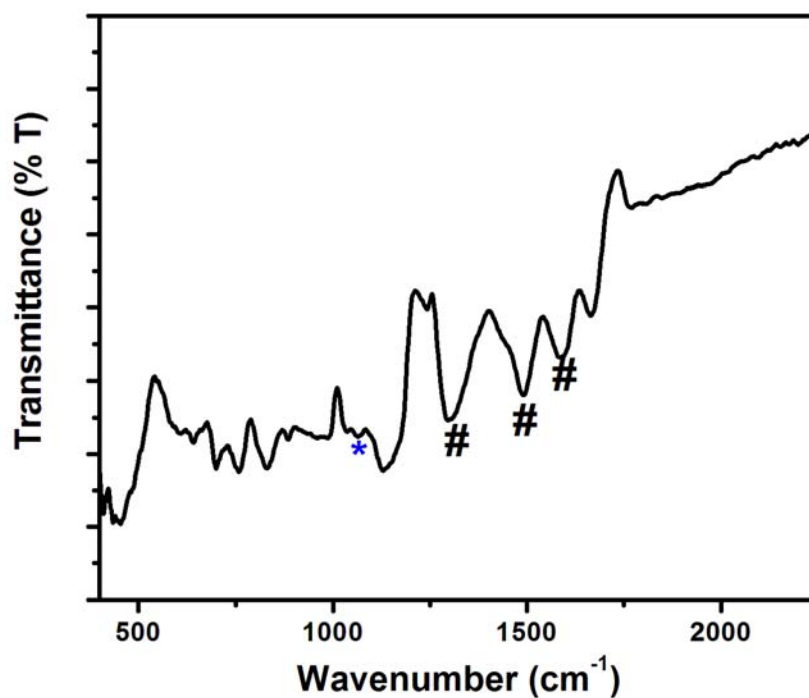
The above FESEM/ TEM images clearly show the formation of PANI- NF's at 60⁰C. The PANI anchored SVO nanospheres also found the formation of NFs synthesized at 80⁰C.

SI₃ XRD of PANI-NF's synthesized at 60⁰C



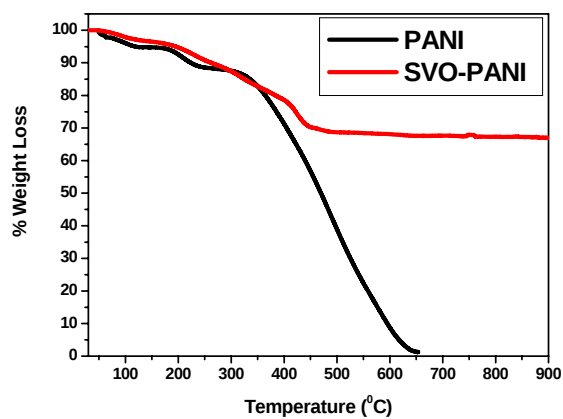
The formation of NF's is characterised by XRD where $2\theta = 6.5^{\circ}$ is the characteristic of PANI-NF's. The peaks at 19.0° and 20.0° indicate the nanofibers of PANI which are partially crystalline.^{24a}

SI₄.FTIR Spectra of layer of PANI - SVO nanospheres NC recorded at 400- 2500cm⁻¹

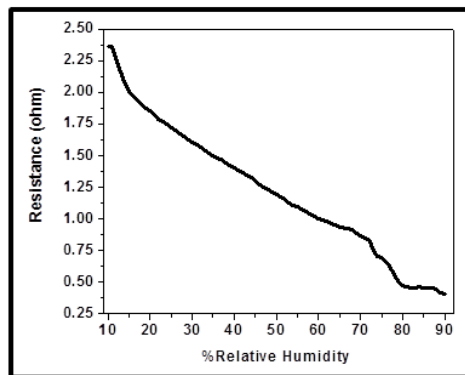


The role of silverous material in PANI layer SVO nanospheres is important where the interaction of inorganic and organic phase elucidated with the help of FTIR shown by (*).

S₅ (a-b) TGA of PANI-NF's and thin layer of PANI - SVO NC.

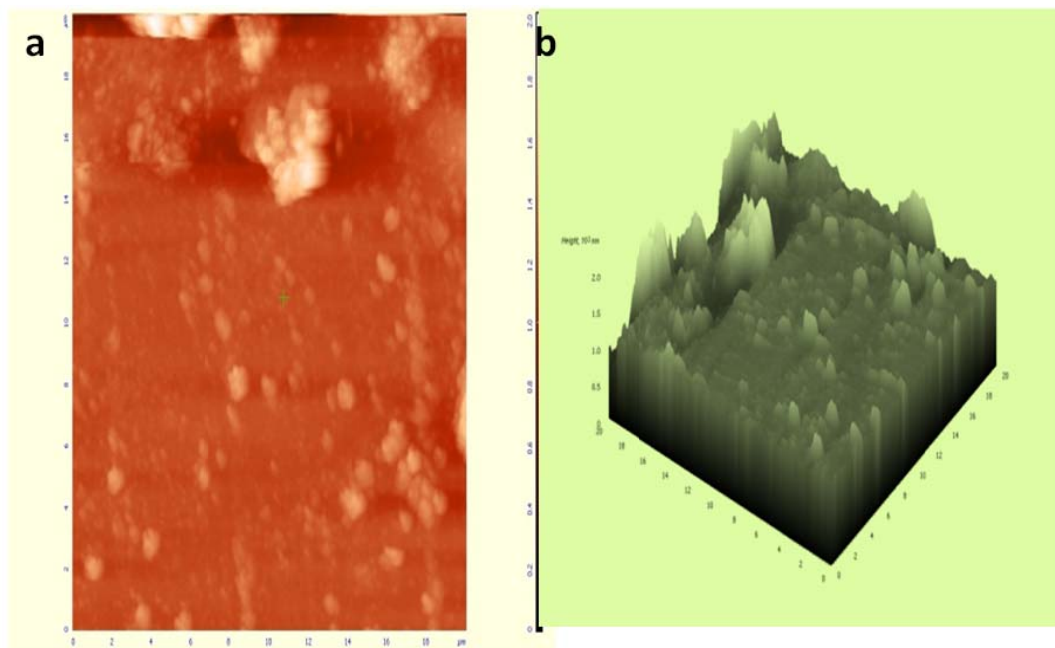


SI₆. Characteristic response of PANI anchored SVO nanocomposites for humidity



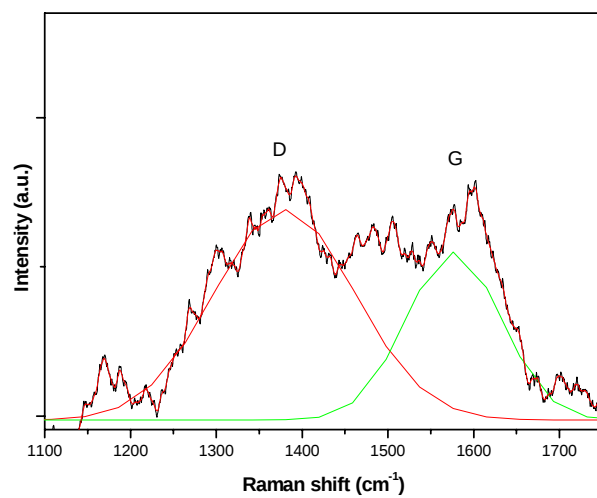
SI-7 PANI - SVO NCs synthesized at 80°C

The height profile of ultrathin layer (2nm) of PANI accompanying SVO supported by XRD and Raman spectra as shown in SI-8.

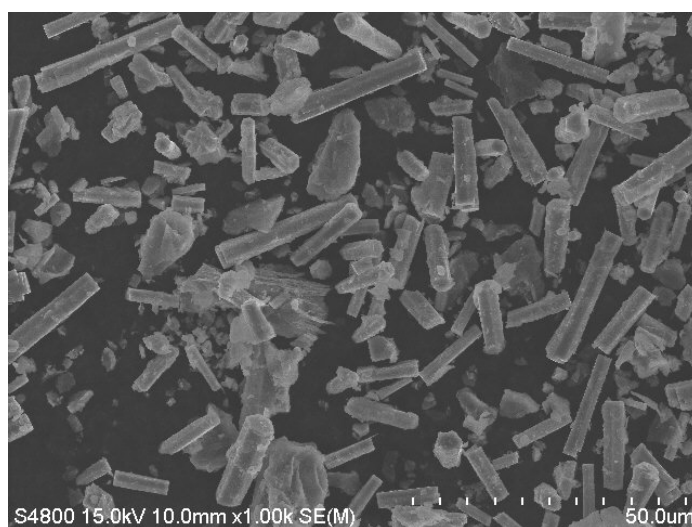


SI-8

Raman spectra of growth of SVO in thin layer of PANI sample have Raman peaks that are similar in shape and position. The peak intensity ratio $I(D)/I(G)$ is ~ 1 . This indicates that the more size of SVO nanospheres, in-plane sp^2 domain is partially ordered. However, in polymerization the ratio is ~ 1 , which suggests that the defect concentration is very low and that the aromatic structures recovered during the post polymerization and hence thin layer of PANI.^{22a}



SI-9 Morphology of SVO Nano / microrods



SI-10 XRD of SVO Nano / microrods

