

Supporting Information

Dwindling the Re-stacking by Simultaneous Exfoliation of Boron Nitride and Decoration of α -Fe₂O₃ Nanoparticles using Solvothermal Route

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Structural characterization:

The formation of α -Fe₂O₃ and BN-Fe₂O₃ composite was primarily characterized by XRD analysis. XRD patterns were recorded using BRUKER D8 ADVANCE X-ray Diffractometer with Cu K α radiation ($\lambda = 1.5418 \text{ \AA}$) from 10-80° at 0.02° step and a count time of 0.2s. FT-IR and Raman analysis was performed using TENSOR 27 spectrometer (Bruker) and laser Raman system (RENISHAW Invia laser Raman microscope) with He-Ne laser (wavelength $\lambda = 633 \text{ nm}$), respectively. Diffuse reflectance spectra (DRS) was carried out using a UV-VIS-NIR double beam spectrophotometer (VARIAN, Cary 5000) equipped with a diffuse reflectance attachment. The decoration of α -Fe₂O₃ nanoparticles over E-BN nanosheets were evaluated by scanning electron microscopy (SEM) and high resolution transmission electron microscopy (HR-TEM) analysis. SEM and HR-TEM images were taken at different locations/magnifications using Hitachi S-3000H (Japan) with an accelerating voltage of 0.3–30 kV and Tecnai G² 20 working at an accelerating voltage of 200 kV, respectively.

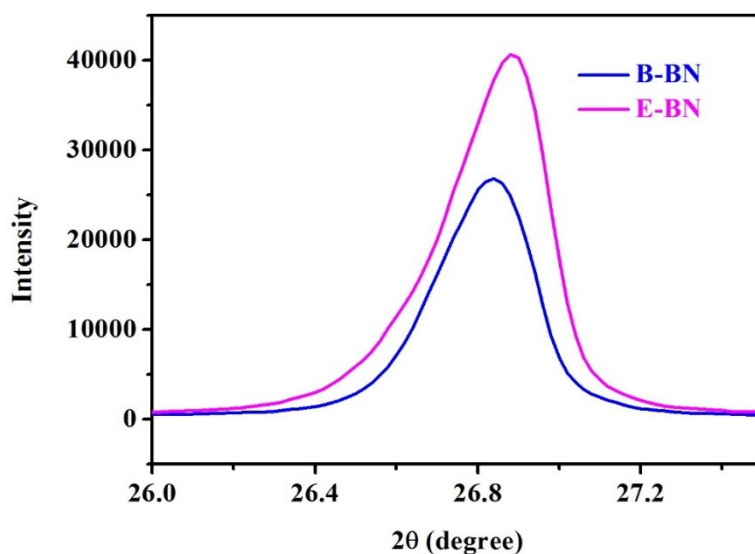


Fig. S1 Enlarged XRD patterns of B-BN and E-BN.

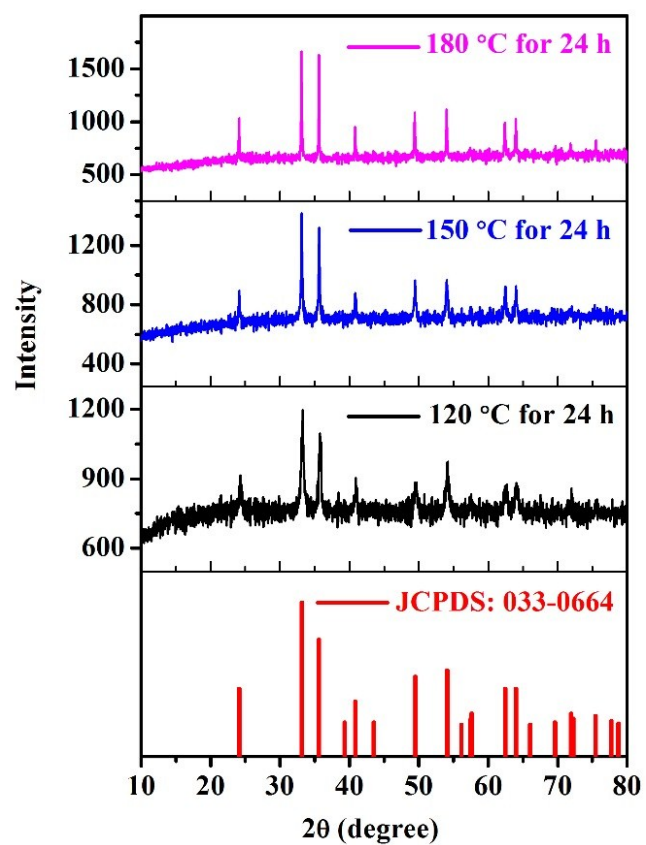


Fig. S2 XRD patterns of α -Fe₂O₃ nanoparticles obtained by different reaction temperatures and the standard reference pattern of α -Fe₂O₃.

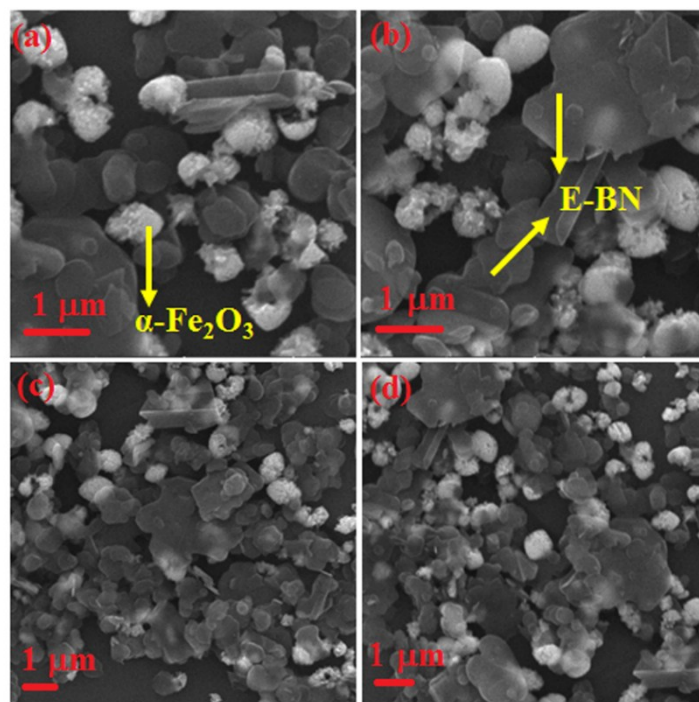


Fig. S3 SEM images of BN-Fe₂O₃ composite (1:1 wt%).

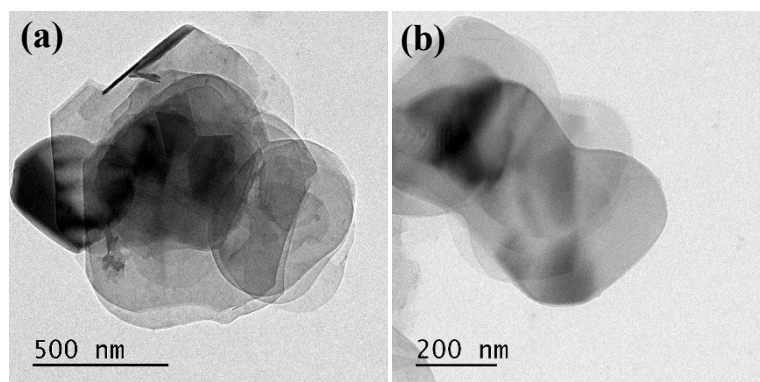


Fig. S4 HR-TEM images of E-BN nanosheets at different magnifications.

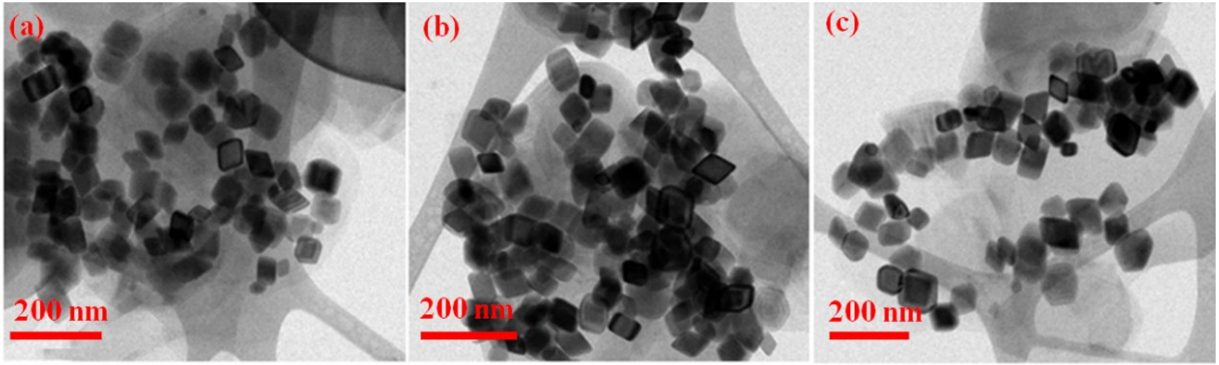


Fig. S5 HR-TEM images of BN-Fe₂O₃ composite.

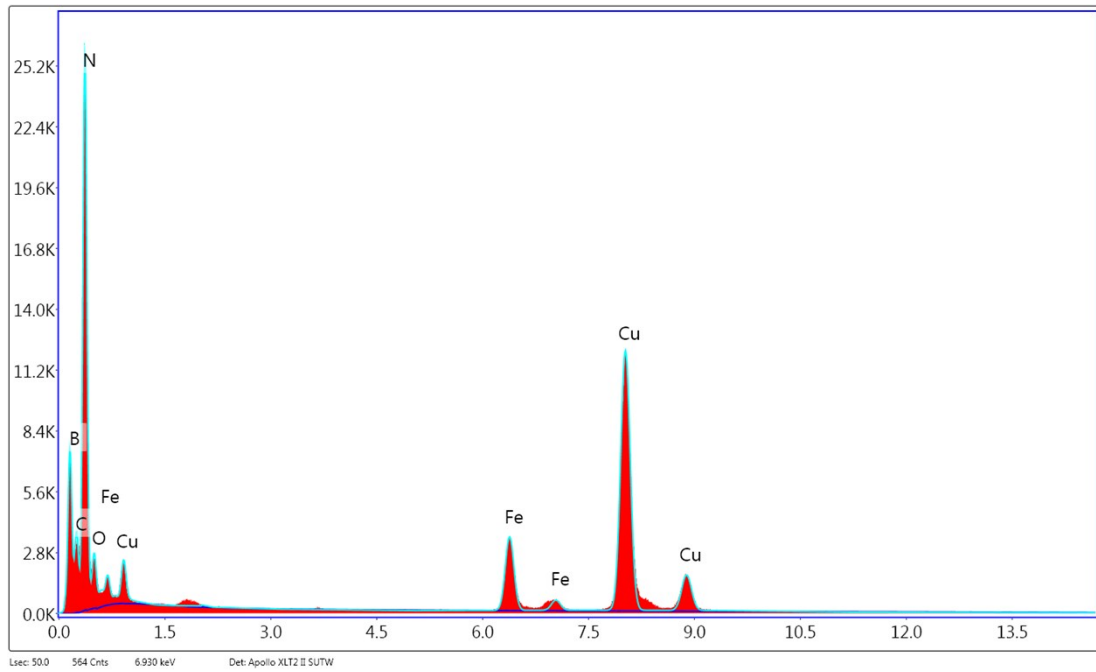


Fig. S6 TEM-EDX analysis of BN-Fe₂O₃ composite.

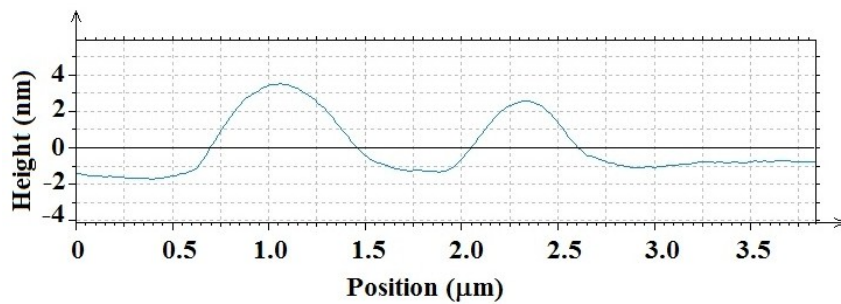
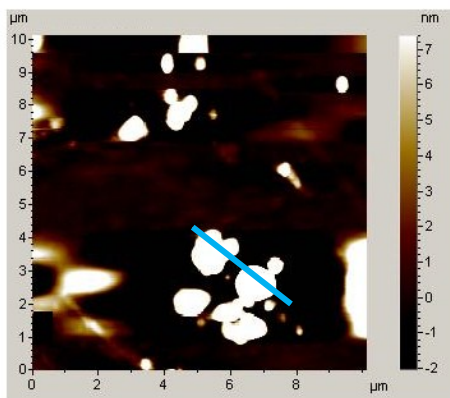


Fig. S7 AFM image of E-BN nanosheets.