Electronic Supplementary Information

Title: Efficient Fe₃O₄ Nanoparticle Catalysts for Depolymerization of Polyethylene Terephthalate

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Fig. S1. TEM images of Fe_3O_4 prepared by various synthesis methods. (a) $MnFe_2O_4$ -CP, (b) Fe_3O_4 -CP, (c) $CoFe_2O_4$ -CP, (d) $NiFe_2O_4$ -CP, and (e) $ZnFe_2O_4$ -CP.



Fig. S2. TEM images of Fe_3O_4 prepared by various synthesis methods. (a) MnFe₂O₄-H-T, (b) Fe_3O_4 -H-T, (c) $CoFe_2O_4$ -H-T, (d) NiFe₂O₄-H-T, and (e) ZnFe₂O₄-H-T.



Fig. S3. XRD patterns of MFe_2O_4 , M=Mn, Fe, Co, Ni, and Zn (a) catalysts prepared by the H method using Triton X-100, and (b) catalysts prepared by the CP method. Asterisks indicate peaks corresponding to metallic Ni (ICSD number 53807).



Fig. S4. Effect of surfactant removal on Fe_3O_4 -TD-s NPs. (a) Photographs of waxy (left) and fine powder (right) forms of Fe_3O_4 -TD-s NPs before and after the washing process. (b) TEM images, and (c) TGA curves of as-synthesized (left) and heavily washed (left) Fe_3O_4 -TD-s NPs. (d) Changes in PET conversion and BHET yield.



Fig. S5. Catalytic performance of PET glycolysis over mixed ferrite (MFe₂O₄-H-T) NPs.



Fig. S6. Temperature-programmed desorption (TPD) profiles by ammonia for MFe₂O₄-H-T NPs.



Fig. S7. Effect of reaction time in PET glycolysis. Reaction conditions: PET 1 g, EG 3 mL, catalyst 0.01 g, and reaction temperature 195°C.



Fig. S8. Product distribution of BHET as a function of glycolysis reaction time by HPLC analysis.



Fig. S9. DSC curves of three kinds of PETs: Commercial PET, r-PET-w/o-A, and r-PET-w/o-A.



Fig. S10. Reusability test of Fe₃O₄-CP NPs in PET glycolysis.



Fig. S11. (a,b) TEM images and (c) XRD patterns of the fresh and spent Fe_3O_4 -CP NP catalysts after reusability experiments.