

Supporting Information

Uniform Epitaxial Growth of Pt on Fe₃O₄ Nanoparticles; Synergetic Enhancement to Pt Activity for the Oxygen Reduction Reaction

*Donald A. Robinson and Keith J. Stevenson**

Department of Chemistry (1 University Station A5300)

The University of Texas at Austin, Austin, Texas 78712, United States.

*stevenson@cm.utexas.edu

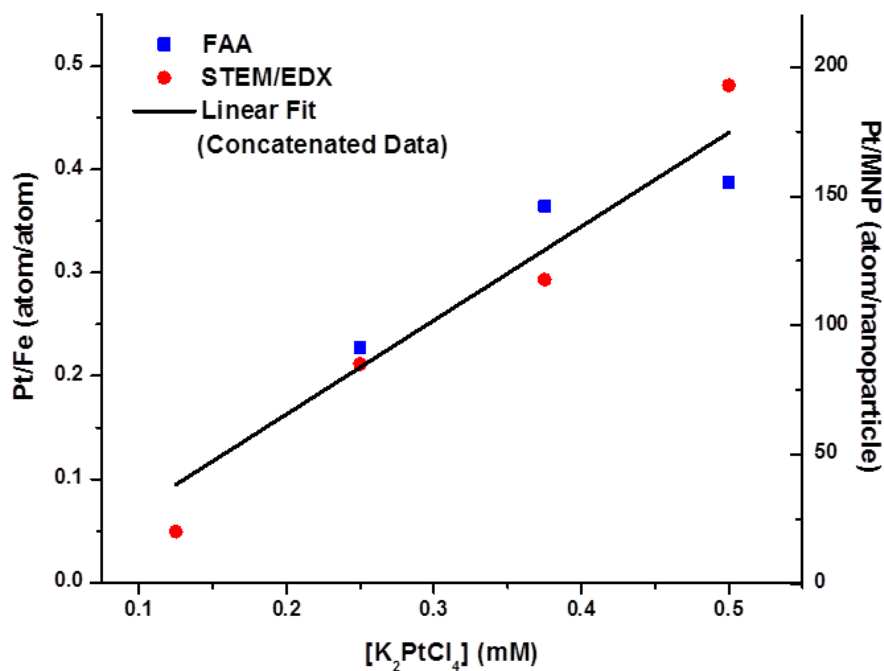


Fig. 1 Relationship between K_2PtCl_4 concentration and resulting composition of PtBPC-MHNs, plotted using both FAA and STEM/EDX data with y-axis scales for both Pt/Fe atomic ratio and Pt/MNP atom/nanoparticle ratio.

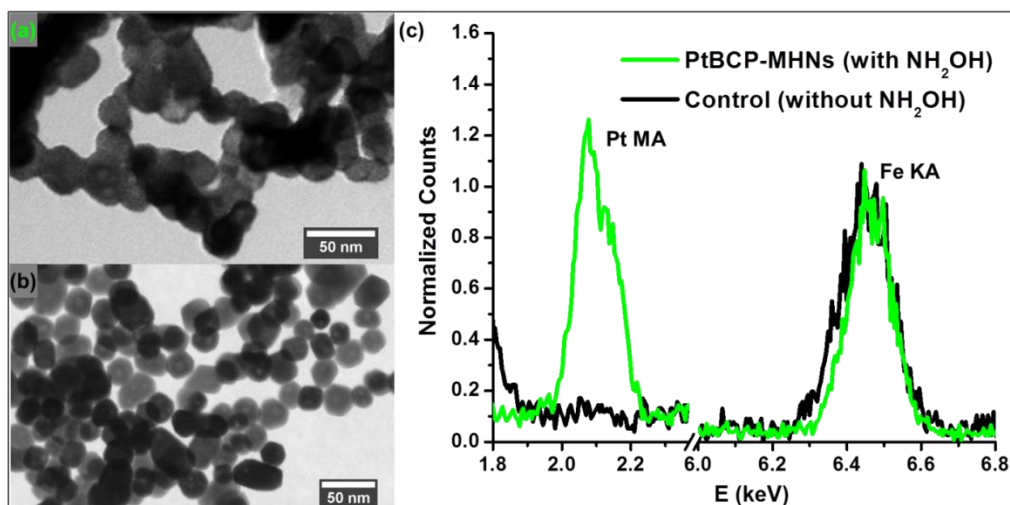


Fig. 2 STEM of nanoparticles resulting from (a) PtBCP-MHN synthesis and (b) control synthesis without NH_2OH . (c) Corresponding EDX spectra for both species, spectra normalized by Fe KA peak maximum.

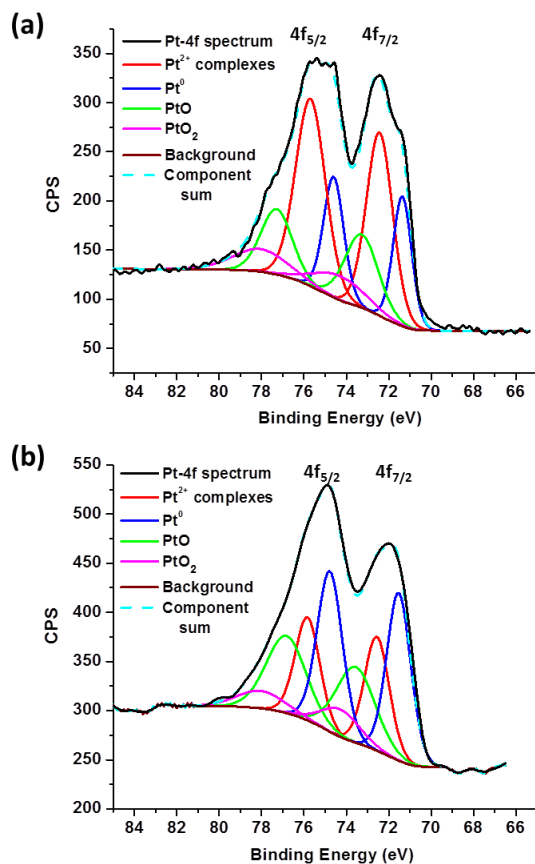


Fig. 3 Deconvoluted Pt-4f XPS spectra corresponding to PtMHN/C samples after thermal treatment in air at a) 190°C and b) 240°C.

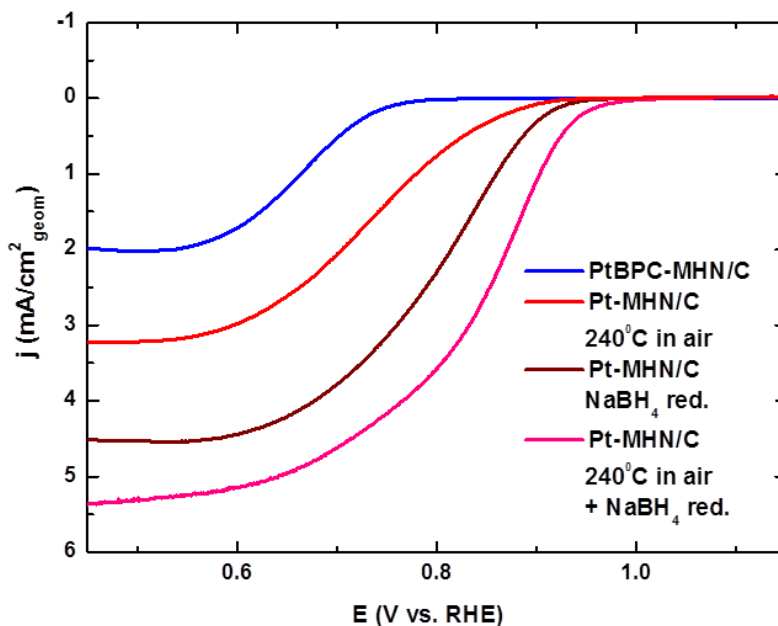


Fig. 4 Rotating disk voltammograms (cathodic scan, 1600 rpm, 5 mV/s, O₂-sat. 0.10 M KOH) for AT-MNP/C, PtBPC-MHN/C, and three different types of Pt-MHN/C species, synthesized under different conditions as labeled in the legend.

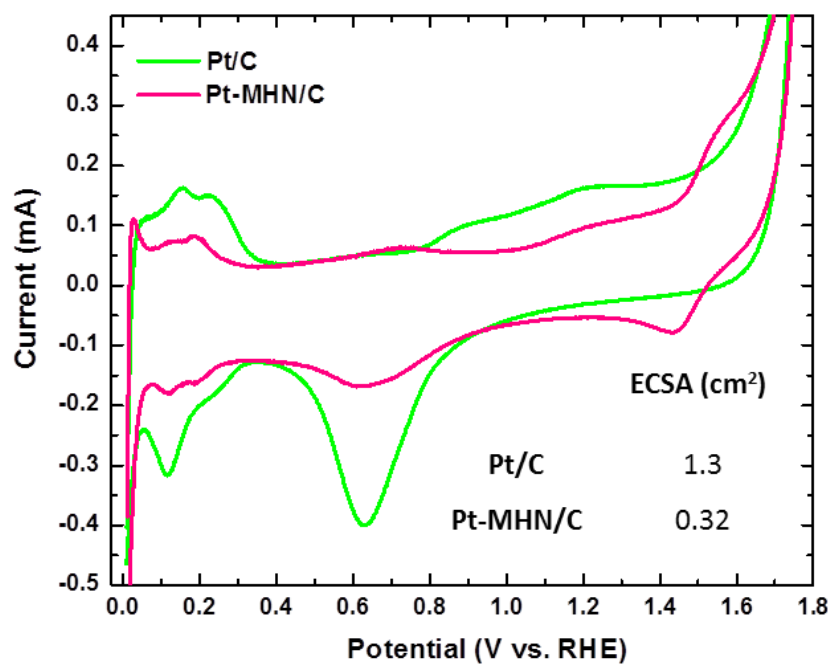


Fig. 5 Cyclic voltammograms (100 mV/s, argon-purged 0.10 M HClO₄) for Pt/C commercial catalyst and Pt-MHN/C. Inset showing ECSA values for each catalyst film.

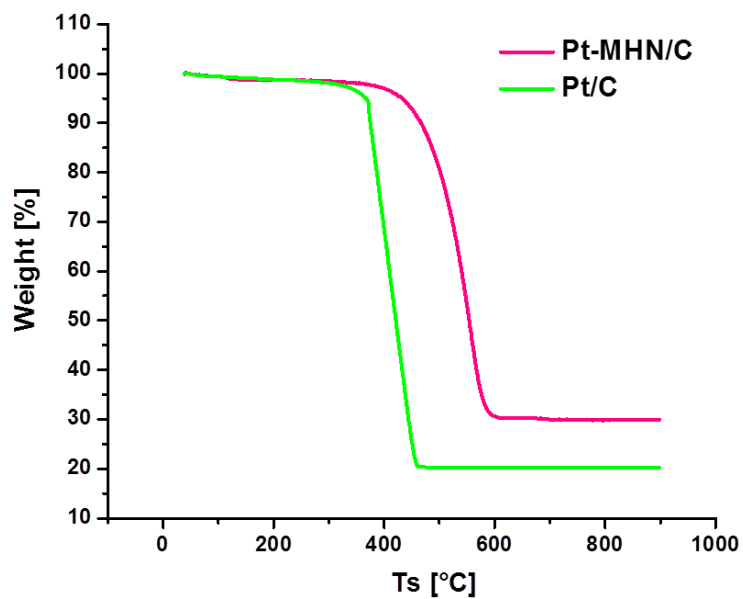


Fig. 6 Thermogravimetric analysis for Pt-MHN/C and Pt/C commercial catalyst.

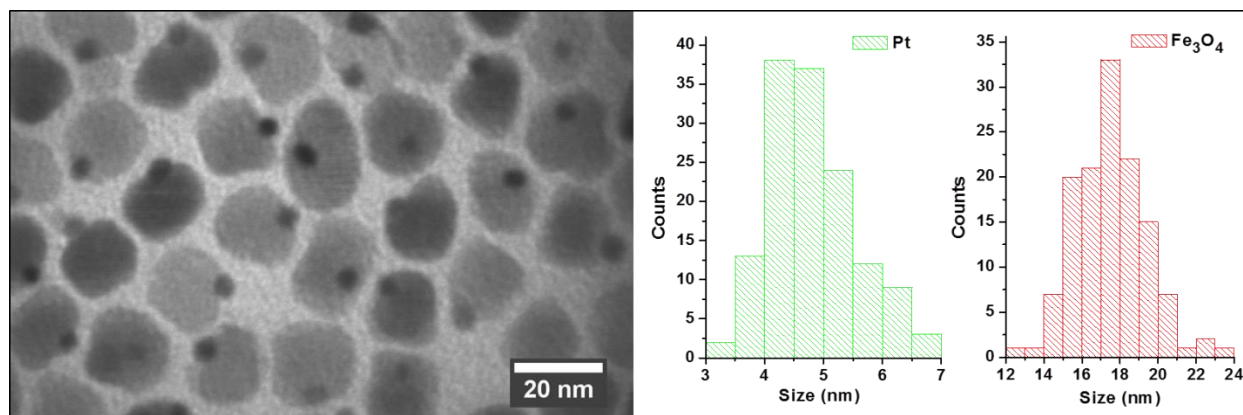


Fig. 7 STEM image and size distribution histograms for Pt-Fe₃O₄ heterodimers (Pt-MHDs).