

Fig. A. UV-Vis absorption spectrum of PB nanoparticles in solution.

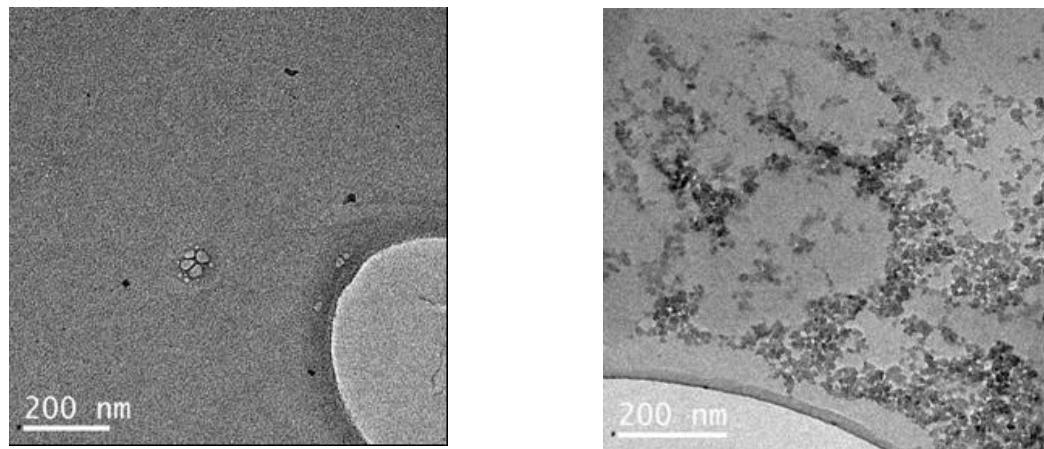


Fig. B. TEM images for PB nanoparticles using $\beta\text{-CD}/\text{Fe}^{2+}$ ratio of 20 (left) and 1.5 (right).

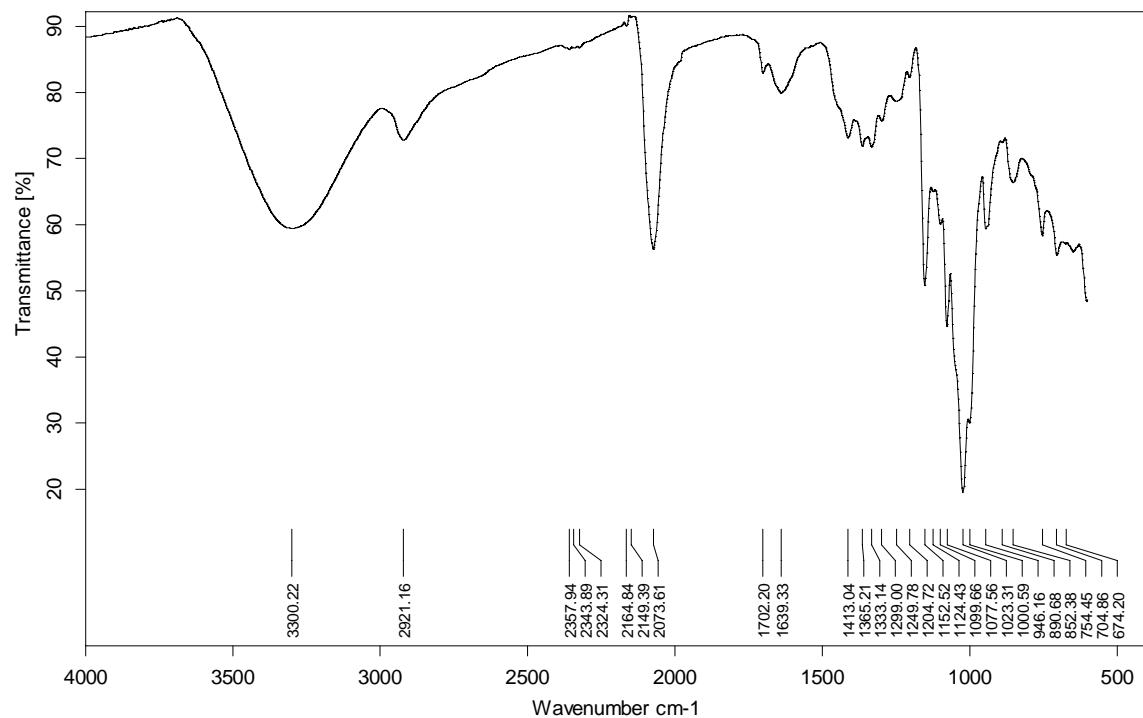


Fig. C. FTIR spectrum of PB-CD using β -CD/ Fe^{2+} stoichiometry ratio of 10.

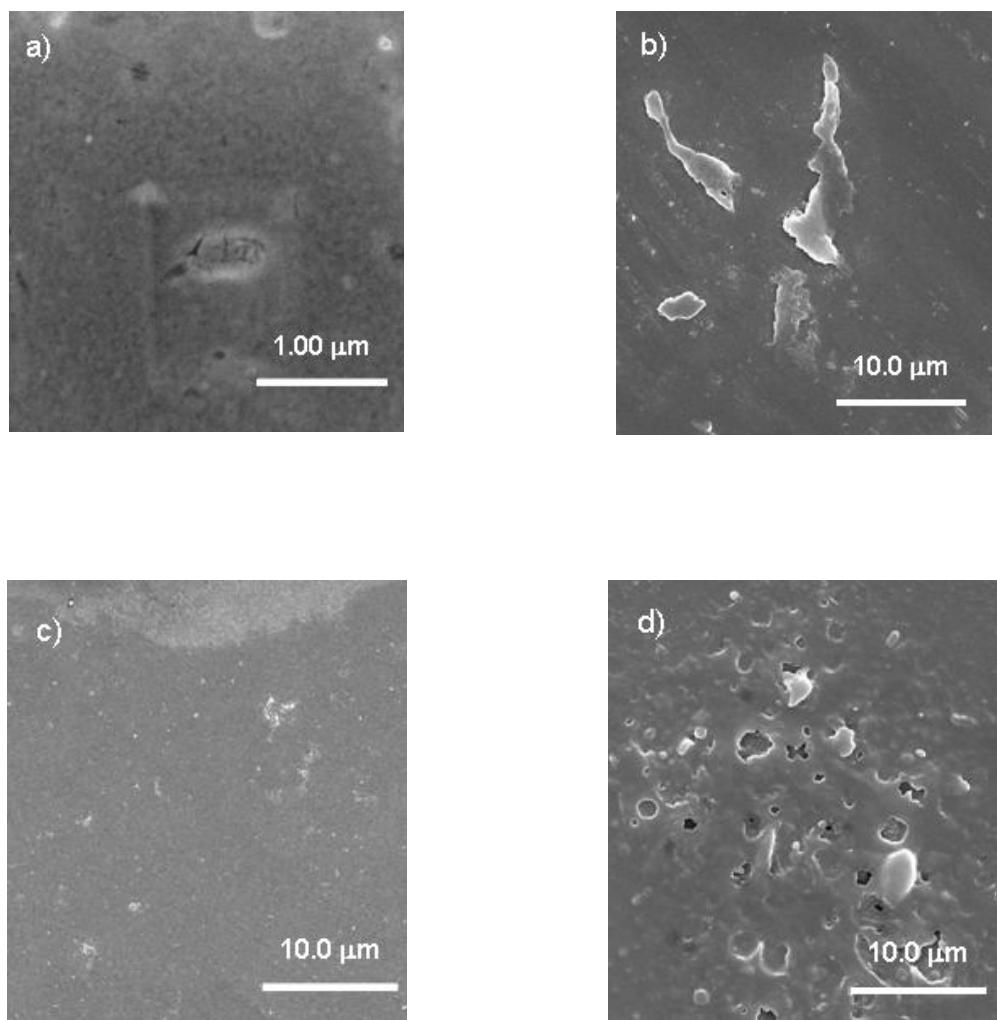


Fig. D. SEM images for: a) PAH, b) PB and c) PB-CD monolayers and d) $\{\text{PAH}/\text{PB-CD}\}_1$ deposited onto ITO substrates.

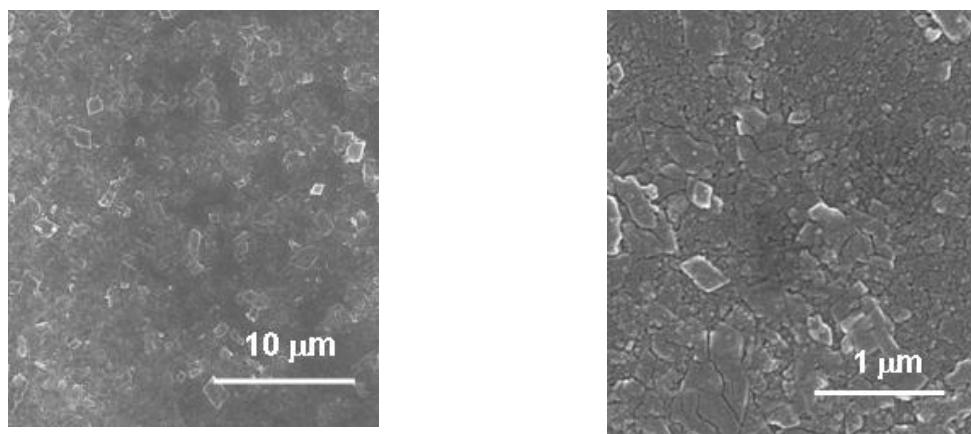


Fig. E. SEM images for $\{\text{PAH}/\text{PB-CD}\}_2$ (left) and $\{\text{PAH}/\text{PB-CD}\}_4$ (right) deposited onto ITO substrates.

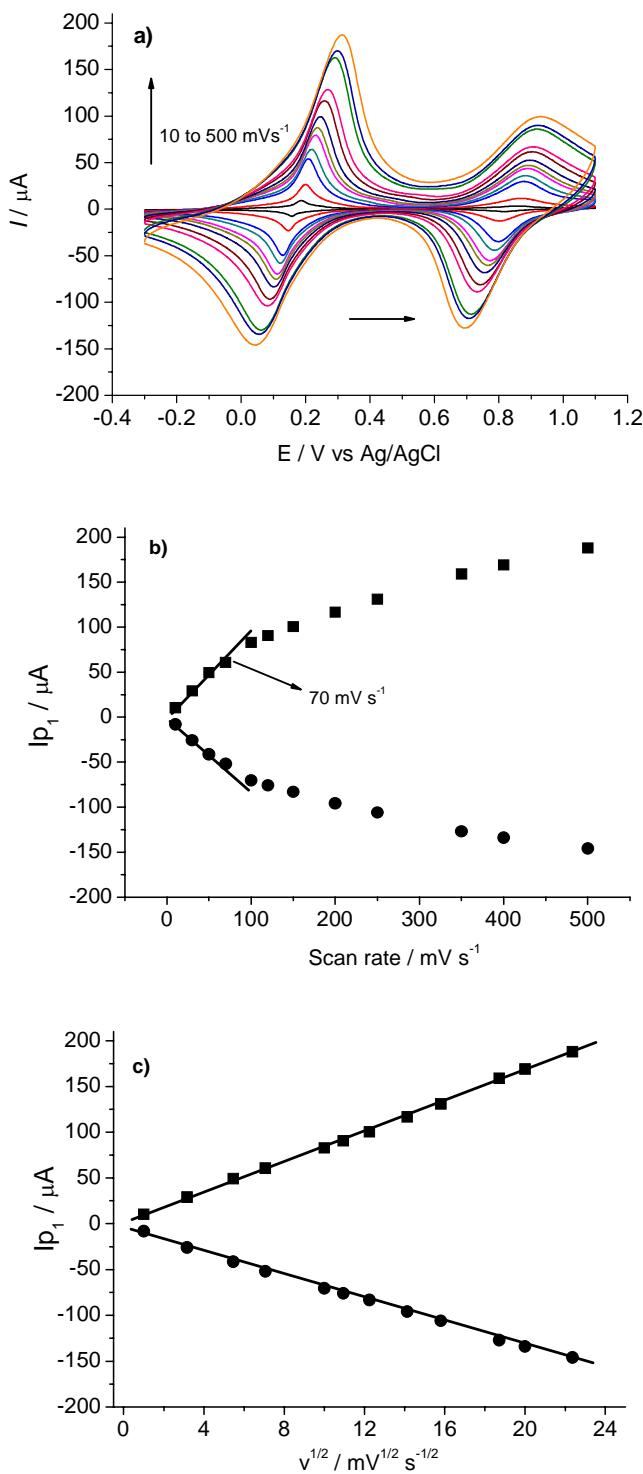


Fig. F. Cyclic voltammograms for: **a)** ITO covered with 3-bilayers of {PAH/PB-CD} in 0.2 mol L⁻¹ KCl solution at scan rate of 10, 30, 50, 70, 100, 120, 150, 200, 250, 350, 400 and 500; **b)** Influence of scan rate (10 to 500 mV s⁻¹) on oxidation and reduction peak currents for ITO-{PAH/PB-CD}₃

electrode; c) Plot of the anodic and cathodic peak currents from E_{p1} as function of the square root of scan rate.

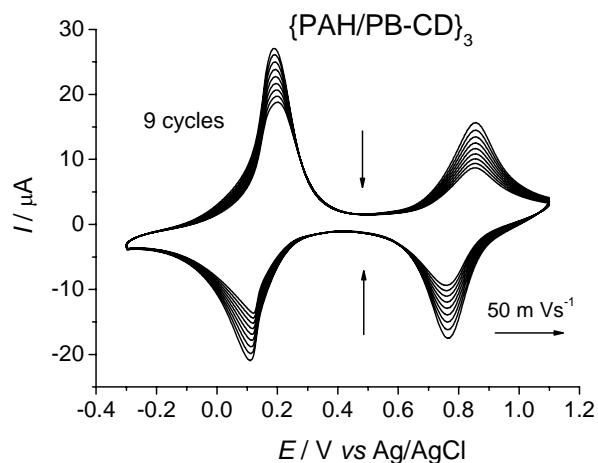


Fig. G. Continuous cyclic voltammograms (9 cycles) for ITO- $\{\text{PAH/PB-CD}\}_3$ modified electrode in 0.2 mol L⁻¹ KCl solution. Scan rate = 50 mV s⁻¹, T = 25 °C.

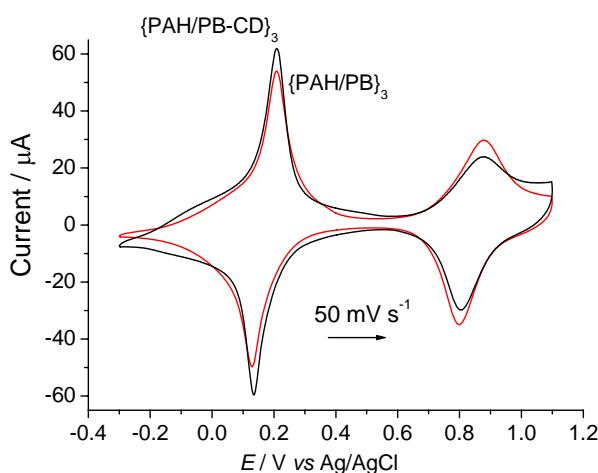


Fig. H. Cyclic voltammograms of 3-bilayers for PAH/PB-CD and PAH/PB systems in 0.2 mol L⁻¹ KCl solution. Scan rate = 50 mV s⁻¹, T = 25 °C.