

Mohammad Ali Kamyabi* and Mohammad Ali Shafiee

Department of Chemistry, Zanjan University, PO Box 45195-313, Zanjan, Islamic Republic of Iran

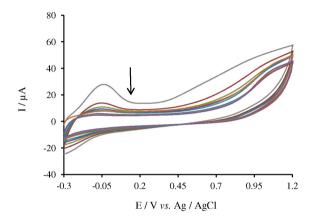


Figure S1. Cyclic voltammograms recorded of poly(2,6-DAP) films on the MWNTs/GCE electrode in 0.01 mol L⁻¹ NaOH solution (alkaline treatment) between -0.3 and 1.2 V vs. Ag / AgCl, potential scan rate 50 mV s⁻¹.

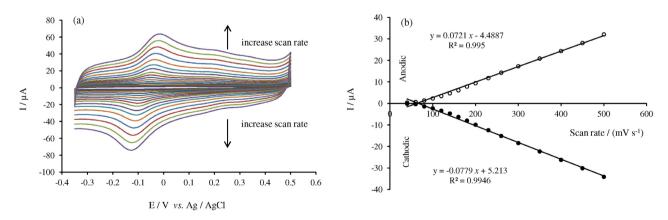


Figure S2. (a) Cyclic voltammograms of poly(2,6-DAP)/MWNTs/GCE in the 0.1 mol L⁻¹ PBS pH 8.0 with varying scan rate, scan rates (mV s⁻¹) of 40, 60, 80, 100, 120, 140, 160, 180, 200, 230, 260, 300, 350, 400, 450 and 500, and (b) plot of cathodic and anodic current *vs.* scan rate (v).

*e-mail: makamyabi@gmail.com

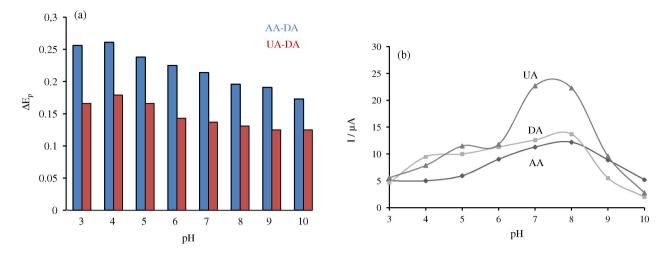


Figure S3. (a) Anodic peak potential differences (E_{pa}) between AA-DA and DA-UA of ascorbic acid (0.5 mmol L⁻¹), dopamine (100 µmol L⁻¹) and uric acid (0.5 mmol L⁻¹) and (b) anodic peak current of AA, DA and UA at the poly(2,6-DAP)/MWNTs/GCE in 0.1 mol L⁻¹ buffer solution with different pH values. Scan rate was 100 mV s⁻¹.

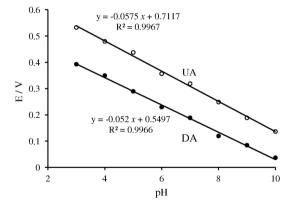


Figure S4. Dependence of the oxidation peak potentials of 100 μ mol L⁻¹ dopamine and 0.5 mmol L⁻¹ uric acid on pH at the poly(2,6-DAP)/MWNT modified GC electrode in 0.1 mol L⁻¹ buffer solution with different pH values. Scan rate was 100 mV s⁻¹.

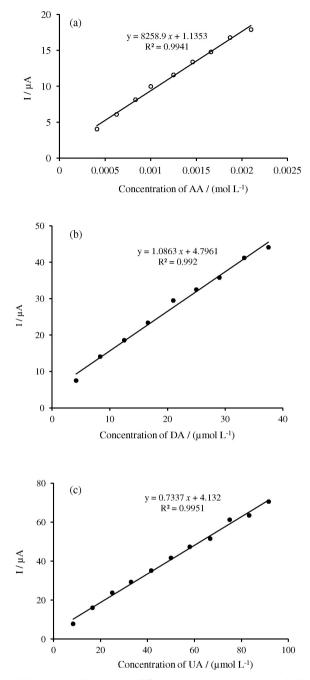


Figure S5. Calibration curve for (a) AA, (b) DA and (c) UA based on differential pulse voltammograms, in PBS pH 8.0, the scan rate was 100 mV s⁻¹.