



Fig.S1. Cyclic voltammograms of the electro-polymerization of Aspartic acid on TOHS/MWCNTs/CPE surface in 0.1 M PBS (pH 7.0) containing 0.05 M Aspartic acid in the potential range of -0.2 to 2.0 V at 100 mV·s⁻¹ for 15 cycles.



Fig. S2. The FTIR spectra of aspartic acid (a) and P-ASP film (b).



Fig.S3. The variations of differential pulse anodic peak currents of 150 μ M of LD and 50 μ M of ACT with respect to accumulation time.



Fig.S4. Cyclic voltammograms 100 μ M LD and 50 μ M ACT at different scan rates (from inner to outer) 0.01, 0.02, 0.03, 0.04, 0.05, 0.06, 0.07, 0.08, 0.09,0.1, 0.12, 0.14, 0.16, 0.18, 0.2, 0.24, 0.28, 0.32, 0.36, 0.4 V.s⁻¹. Insets: (a) peak separations (ΔE_p) for ACT as a function of log (v), (b) peak potential for LD as a function of ln (v) and (c) peak potential for LD as a function of ln (v).