Supplementary Material

Simultaneous determination of dopamine and uric acid in presence of high concentration of ascorbic acid using cetyltrimethylammonium bromide– polyaniline/activated charcoal composite

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Electrodes	Dopamine				Uric acid			
	$I_{\rm pa}/\mu{ m A}$	$I_{\rm pc}/\mu{ m A}$	$E_{\rm pa}/{ m V}$	$E_{\rm pc}/{ m V}$	$I_{ m pa}/\mu{ m A}$	$I_{\rm pc}/\mu{ m A}$	$E_{\rm pa}/{ m V}$	$E_{\rm pc}/{ m V}$
Bare GCE	6.237	-1.048	0.384	-0.078	4.057	_	0.566	_
CTAB-PANI	10.47	-2.048	0.315	-0.083	4.585	_	0.400	-
CTAB-PANI/AC	11.88	-3.991	0.218	-0.149	9.587	-1.414	0.335	0.276

Table S1 Comparison of electrocatalytic parameters obtained at CTAB–PANI/AC with control electrodes. The potentials are referred to Ag/AgCl (saturated KCl) reference electrode.

 I_{pa} = anodic peak current, I_{pc} = cathodic peak current, E_{pa} = anodic peak potential and E_{pc} = cathodic peak potential



Fig. S1 CVs obtained CTAB–PANI/AC/GCE in phosphate buffer (pH 7.0) containing 2 mM DA (a) and 200 μ M UA (b) at different scan rates from (a=20, b=40, c=60, d=80, e=100, f=120, g=140, h=160, i=180, j=200 mV s⁻¹). Insets: Plots between I_{pa} (μ A) vs. scan rate (mV s⁻¹)



Fig. S2 Cyclic voltammograms (a) and differential pulse voltammograms (b) obtained at CTAB– PANI/AC/GCE in phosphate buffer (pH 7.0) containing of different concentrations of AA (0.5, 1.0, 1.5 and 2.0 mM).