The preferential electrocatalytic behaviour of graphite and multiwalled carbon nanotubes on endiol groups and their analytical implications in real domains

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Supplementary data



Figure S-1. SEM micrographs of SPE modified with different carbon materials; not modified (A), graphite (B), MWCNT-B (C) and MWCNT-A (D). Conditions: Amplification 5000x, 25 kV.



Figure S-2. Cyclic voltammograms for standards of ascorbic acid (A), hydroquinone (B) and catechin (C) using different modifiers for SPE; bare (black line), Fe_3O_4 nanoparticles (blue line), NiO nanoparticles (green line) and MWCNT-A (red line). Conditions: Scan rate 100mV/s, step potential 0.01V.

-	SPE	SPE-NiO	$SPE-Fe_2O_3$	SPE-MWCNT-A
Ascorbic acid	+0.41	+0.39	+0.48	+0.03
Hydroquinone	$+0.31^{a}$	$+0.28^{a}$	$+0.38^{a}$	$+0.09^{a}$
	-0.07 ^b	-0.06 ^b	-0.10 ^b	$+0.01^{b}$
Catechin	$+0.33^{a}$	$+0.37^{a}$	$+0.40^{a}$	$+0.20^{a}$
	$+0.04^{b}$	-	$+0.08^{b}$	+0.15 ^b

Table S-3. Redox peak potentials for SPE modified with metal oxide nanoparticles.

^{*a*} anodic sweep ^{*b*} catodic sweep



Figure S-4. XPS of C 1s (A) and O 1s (B) for acid treated MWCNT-A (a), untreated MWCNT-A (b).



Figure S-5. Cyclic voltammograms for standards of 1mM ascorbic acid (A), 1mM hydroquinone (B) and 1mM (+)-catechin (C) using different SPE; bare (black line), electrochemically activated (blue line), untreated MWCNT-A (green line) and acid treated MWCNT-A (red line). Conditions: Scan rate 100mV/s, step potential 0.01V.