Graphene layers modified glassy carbon electrode for the determination of

norepinephrine and theophylline in pharmaceutical formulations

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Fig. S1. Electrochemical reduction of electrostatically assembled GO on ITO/HDA electrode (15 cycles) in PB solution (pH 7.2; 0.2 M) at a scan rate of 100 mV s⁻¹.



Fig. S2. XPS survey spectra obtained for (a) ITO/HDA/GO and (b) ITO/HDA/ERGO substrates.



Fig. S3 (A) Electrochemical reduction of electrostatically assembled GO prepared by (a) Hummer's Method, (b) Modified Hummer's method reported by Tour et al., and (c) Tour's method on GC/HDA electrode in 0.2 M PB solution (pH 7.2) at a scan rate of 100 mV s⁻¹. **(B)** CVs obtained for 0.5 mM NE at ERGO modified electrodes fabricated by the electrochemical reduction of GO which was prepared by (a) Hummer's Method, (b) Modified Hummer's method reported by Tour et al., and (c) Tour's method.



Fig. S4. DPVs obtained for 0.5 mM (A) NE and (B) TP at GC/HDA/ERGO electrode in 0.2 M PB solution of pH solution (a) 3, (b) 4, (c) 5, (d) 6, (e) 7 (f) 8 and (g) 9. **Inset**: Plot of pH versus anodic peak potential of AP.



Fig. S5. CVs obtained for the oxidation of 0.5 mM of **(A)** NE and **(B)** TP at GC/HDA/ERGO electrode in PB solution (pH 6; 0.2 M) at different scan rates (a) 25, (b) 50, (c) 75, (d) 100, (e) 125, (f) 150, (g) 175, (h) 200, (i) 225 and (j) 250 mV s⁻¹. Insets: Plots of current versus square root of scan rate.



Fig. S6. Plots of concentration versus peak current for the simultaneous determination of **(A)** NE and **(B)** TP at GC/HDA/ERGO electrode in PB solution (pH 6; 0.2 M).



Fig. S7. DPVs obtained for the increment of 10 μ M TP in the presence of 500 μ M NE at GC/HDA/ERGO electrode in PB solution (pH 6; 0.2 M). Inset: Plot of concentration of TP versus oxidation current.



Fig. S8. DPVs obtained for 20 μ M NE in the presence of 1 mM AA, 500 μ M UA and 200 μ M TP at GC/HDA/ERGO electrode in 0.2 M PB solution.