

Supporting Information

Table SI-1. PTA temperature program for graphene.

Temperature (°C)	Time (s)	Phase
310	90	100 % He
475	90	100 % He
615	90	100 % He
675	90–300	100 % He
500	90	90% He/10% Ox
550	90	90% He/10% Ox
600	90	90% He/10% Ox
700	90	90% He/10% Ox
775	90	90% He/10% Ox
910	90–300	90% He/10% Ox

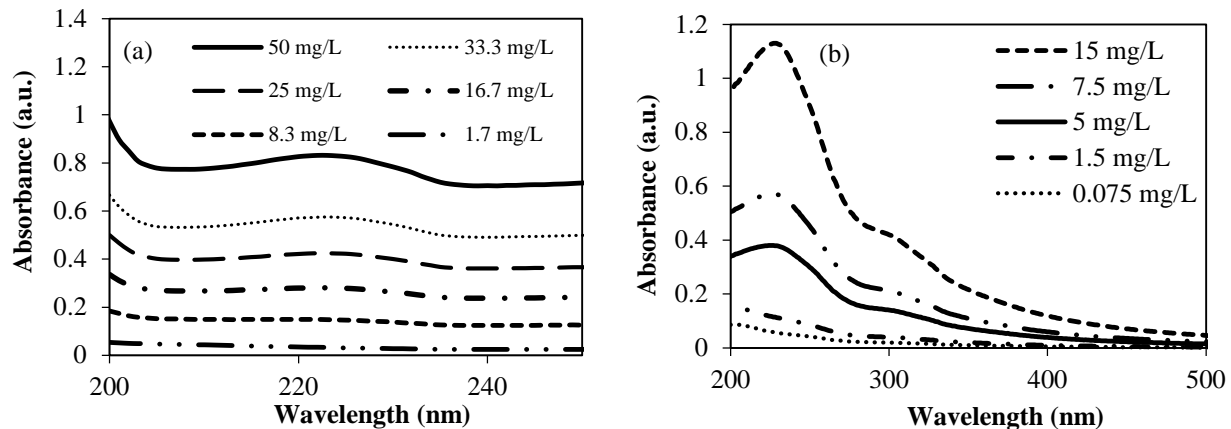


Figure SI-1. UV-Vis absorbance spectra for (a) few-layer graphene (FLG) and (b) graphene oxide (GO).

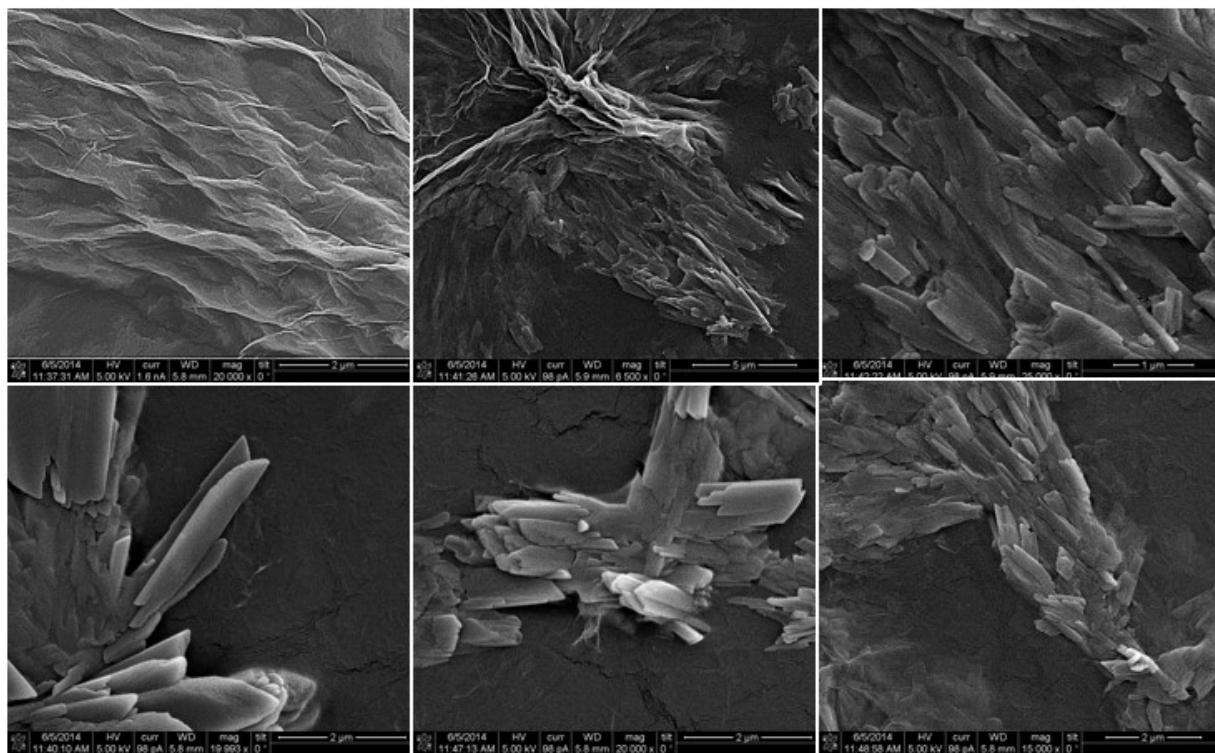


Figure SI-2a. SEM images of GO. The structure of GO ranged from thin, stacked sheets to thick, plate-like particles presumably formed upon drying the aqueous sample.

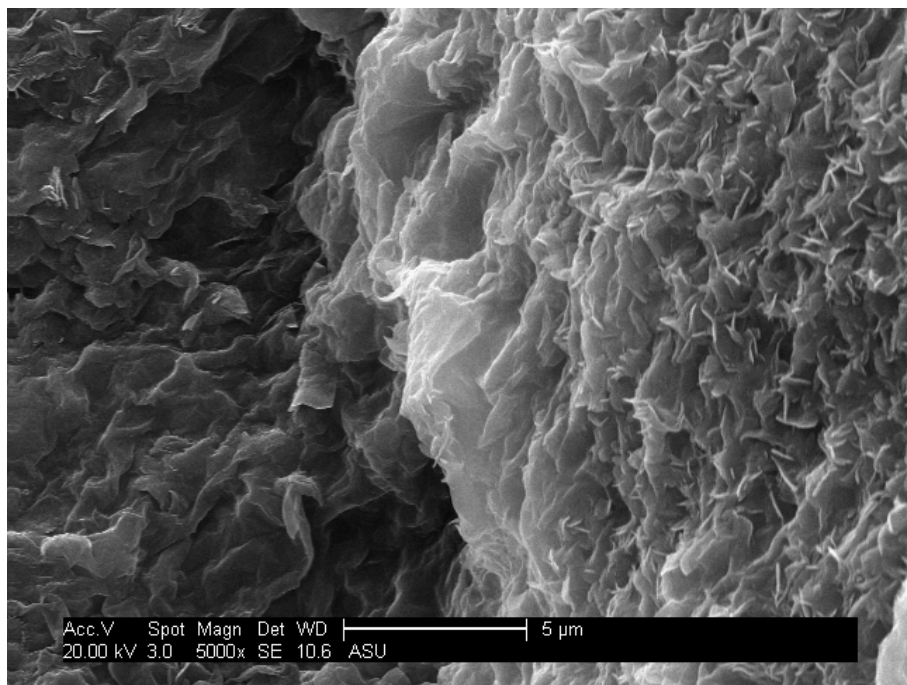


Figure SI-2b. SEM image of RGO.

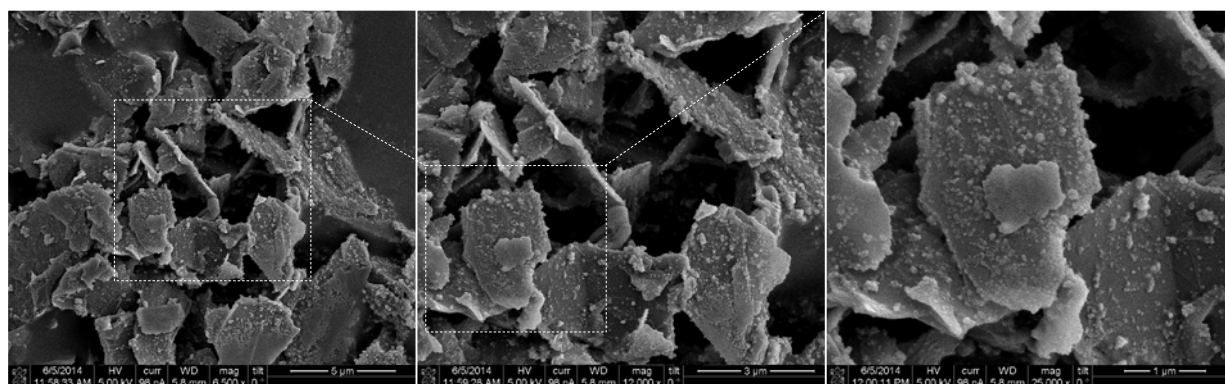


Figure SI-2c. SEM images of FLG. Nanoparticles on surface are from gold sputtering.



Figure SI-3. Image showing graphene sample loaded on a QFF.

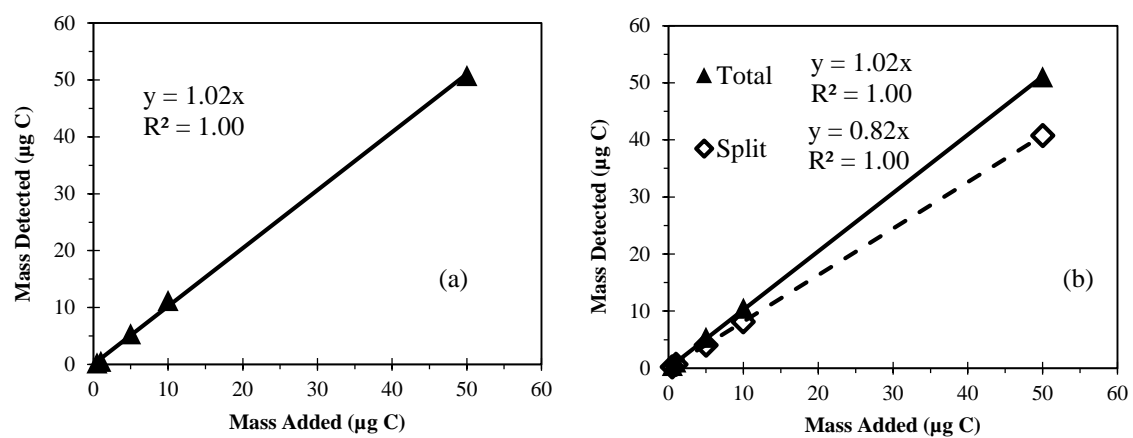


Figure SI-4. Instrument calibration curves for (a) FLG and (b) GO (using two splits). The “Split” values in (b) represent the loss that occurs if accounting only for the carbon evolving during the oxidizing phase.



Figure SI-5. Image showing an example of poor RGO pellet formation in water after centrifugation. 2% NaBH_4 was used for the reduction.

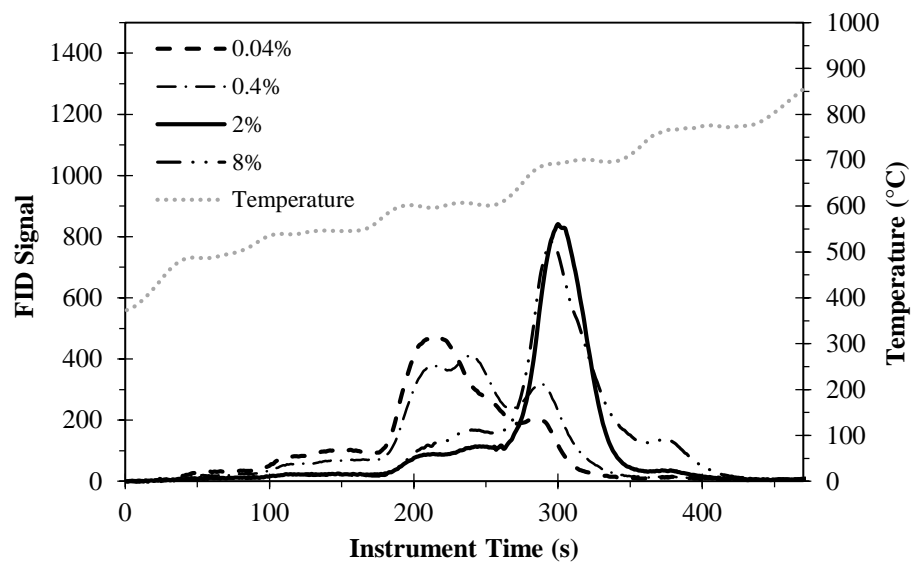


Figure SI-6. PTA thermograms for RGO reduced with different NaBH_4 concentrations in Solvable. Only the oxidizing phase is shown.

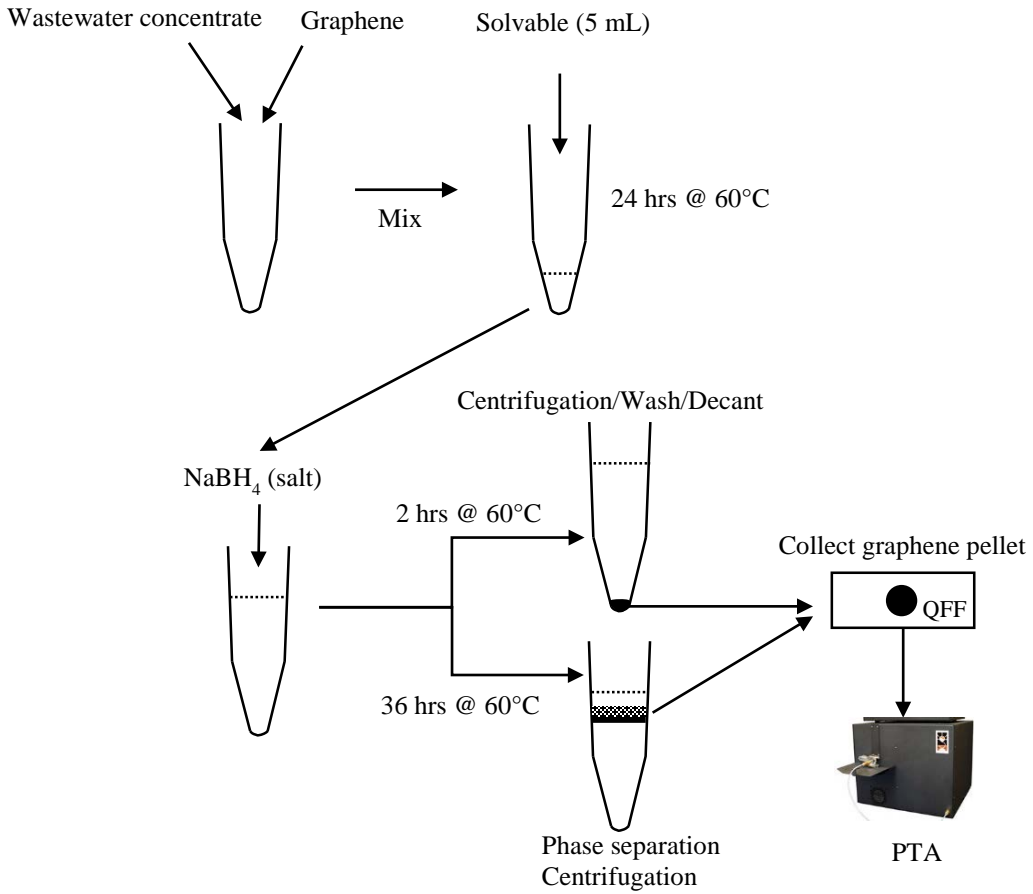


Figure SI-6. Schematic showing the centrifugal and phase separation methods for recovery GO from wastewater.

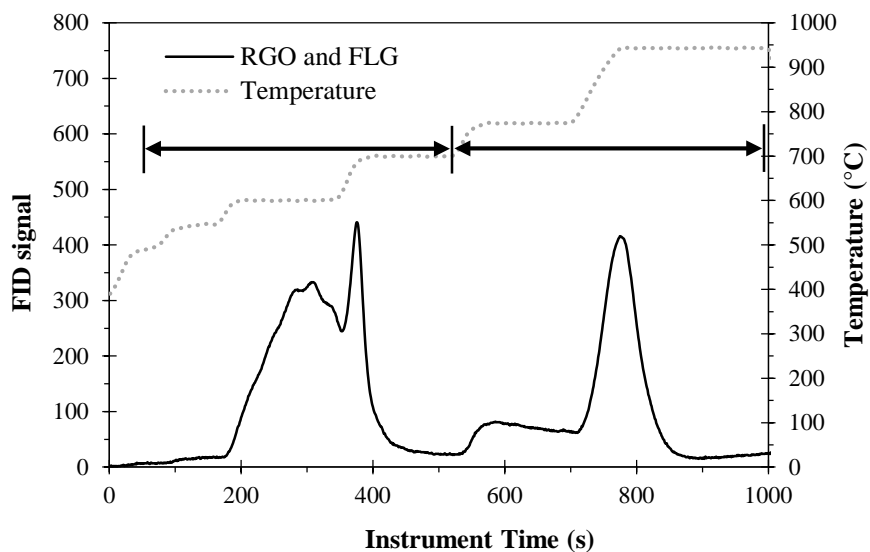


Figure SI-7. PTA thermogram for a mixed sample of GO/RGO and FLG after *in-situ* extraction from a clean matrix (i.e., water, no biomass). GO/RGO was quantified from 500–700°C and FLG was quantified from 700–950°C resulting in recoveries of 85% and 66%, respectively.