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Supplemental Data

Colloidal Stability of Reduced Graphene Oxide Materials Prepared Using Different Reducing Agents

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Procedures used to determine relative hydrophobicity of GO and RGO

Relative hydrophobicity of GO and RGOs was assessed using a hydrocarbon partitioning test with laboratory-grade *n*-dodecane ^{S1}. Samples were prepared by adding 4 ml of a GO suspension to a test tube containing 1 ml of *n*-dodecane. The test tube was vortexed for 2 min, and then left undisturbed for 15 min to allow phase separation. The relative hydrophobicity was assessed as the fraction of GO that partitioned into *n*dodecane from the aqueous phase ^{S2}.

	Artificial surface water	Artificial ground water Concentration (mM)	
Ion type	Concentration (mM)		
Na ⁺	0.23	0.43	
K^+	0.03	0.20	
Ca^{2+}	0.33	0.68	
Ca^{2+} Mg^{2+}	0.15	0.24	
Cl-	0.16	0.64	
NO ₃ -	-	0.62	
HCO ₃ -	0.92	0.45	
SO_4^{2-}	0.07	0.38	

 Table S1 Ion compositions of artificial surface and ground waters.

Table S2 Electrophoretic mobility of GO/RGOs in the absence and presence of SRHA.

	$EPM (m^2V^{-1}s^{-1})$		
	0.5 mM Ca ²⁺	0.5 mM Ca^{2+} with	0.5 mM Ca ²⁺ with
		1 mg/L SRHA	10 mg/L SRHA
GO	-1.24 ± 0.06	-1.27 ± 0.02	-1.34 ± 0.07
VC-RGO	-1.09 ± 0.15	-1.11 ± 0.04	-1.31 ± 0.10
NaBH ₄ -RGO	$\textbf{-}0.99\pm0.01$	-1.00 ± 0.04	-1.11 ± 0.07
N ₂ H ₄ -RGO	-0.91 ± 0.07	-0.98 ± 0.04	-1.11 ± 0.01

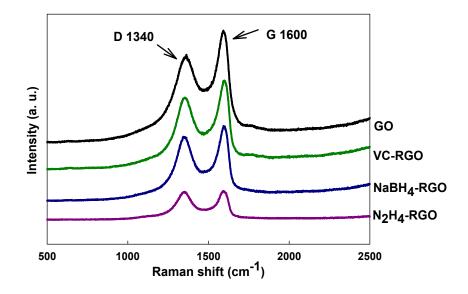
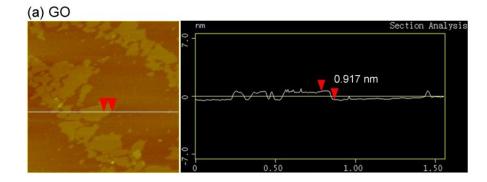
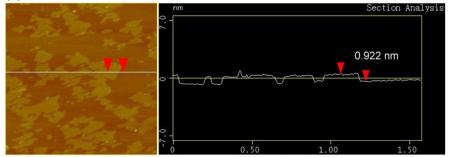


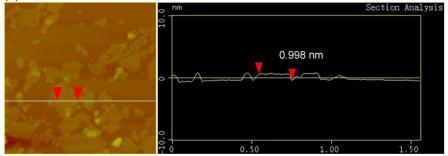
Fig. S1 Raman spectra of GO and reduced GOs. All samples displayed strong Raman peaks at approximate 1340 cm⁻¹ (assigned to the D band) and 1600 cm⁻¹ (G band). The G band is ascribed to the E_{2g} mode of sp² carbon atoms and the D band is usually related to the vibration of carbon atoms with dangling bonds and ascribed to the edge of carbon network ^{S3}.



(b) VC-RGO



(c) NaBH₄-RGO



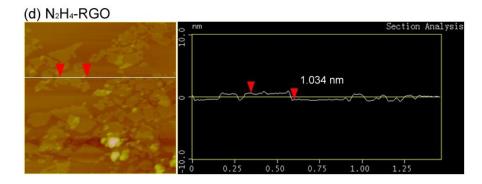
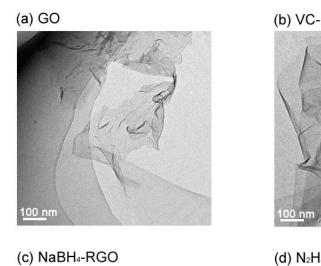
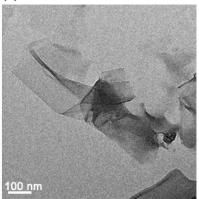
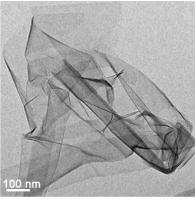


Fig. S2 AFM images and height profiles of graphene oxide (GO) and reduced graphene oxides (VC-RGO, NaBH₄-RGO and N₂H₄-RGO).





(b) VC-RGO



(d) N₂H₄-RGO

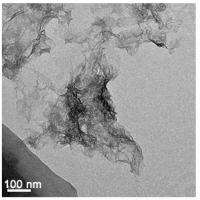


Fig. S3 TEM images of graphene oxide (GO) and reduced graphene oxides (VC-RGO, NaBH₄-RGO and N₂H₄-RGO).

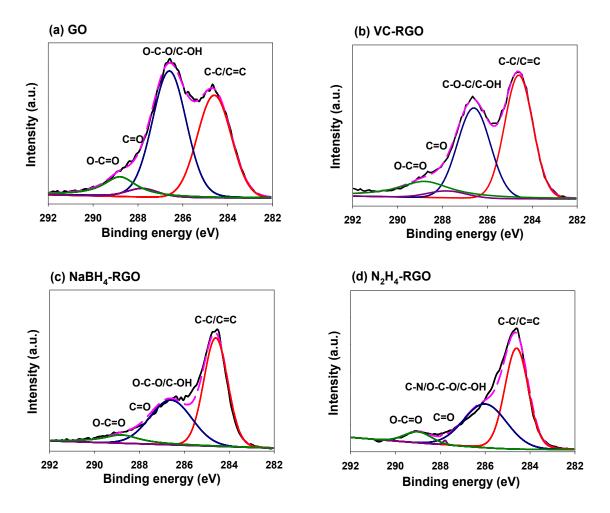


Fig. S4 XPS spectra of graphene oxide (GO) and reduced GOs (VC-RGO, NaBH₄-RGO and N₂H₄-RGO). The peaks with the binding energy of 284.6, 286.6, 287.8 and 288.8 eV are assigned to the carbon atoms in aromatic rings (C-C/C=C), epoxy/hydroxyl (C-O-C/C-OH), carbonyl (C=O) and carboxyl (O-C=O), respectively.

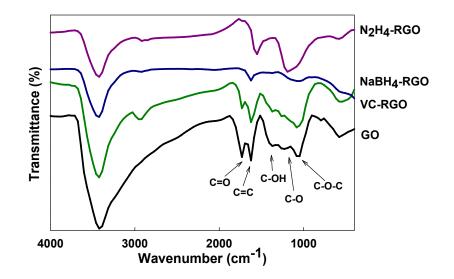


Fig. S5 FTIR spectra of GO and reduced GOs, including VC-RGO, $NaBH_4$ -RGO and N_2H_4 -RGO.

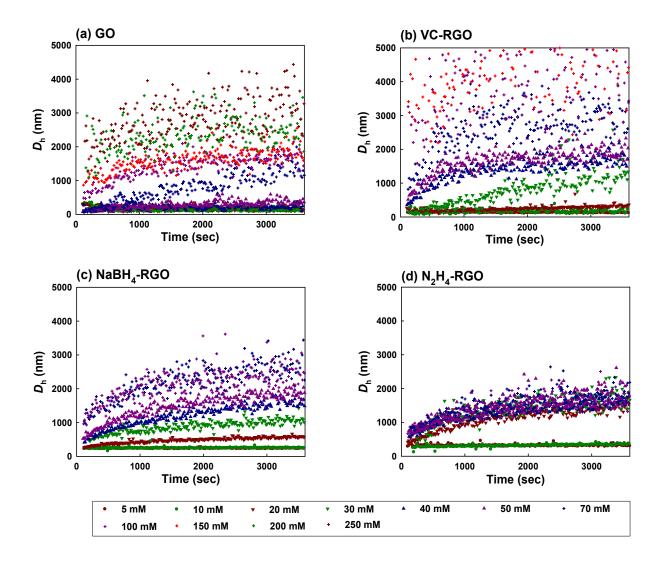


Fig. S6 Aggregation kinetics of GO and reduced GOs (VC-RGO, NaBH₄-RGO and N₂H₄-RGO) as a function of NaCl concentration. Aggregation of GO was studied over a NaCl concentration range of 5 to 250 mM, and that of RGOs was studied over 5 to 150 mM, due to the lower critical coagulation concentrations.

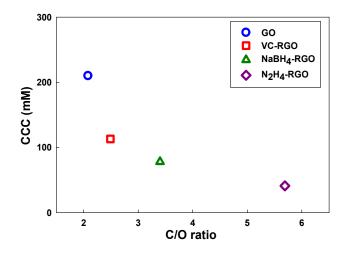


Fig. S7 Correlation of critical coagulation concentration (CCC) values of GO and reduced GOs (VC-RGO, NaBH₄-RGO and N₂H₄-RGO) with C/O ratios of the materials.

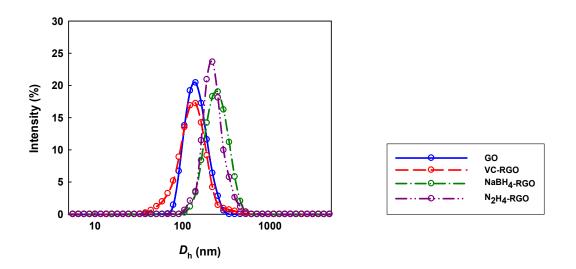


Fig. S8 Particle size distribution of GO and reduced GOs (VC-RGO, NaBH₄-RGO and N_2H_4 -RGO) in deionized water.

References

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