Electronic Supplementary Information:

Redox Chemistry between Graphene Oxide and Mercaptan

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Figure S1. Photos of GO dispersion from control reaction at 80 °C in water in the absence of mercaptan. (A) Original reaction dispersion after heating for 660 min in water; (B) DMF-diluted GO dispersion at different reaction time.



Figure S2. TGA curves of pristine GO and GO after heating in water at 80 °C for 660 min.



Figure S3. Raman spectra of graphite and RGO samples by different mercaptans.



Figure S4. TGA plots of RGO samples reduced by hydroxyethyl mercaptan (HEM) and propyl mercaptan (PM).



Figure S5. Mass spectrum of didodecyl disulfide detected by GC of the reaction mixture.



Figure S6. Mass spectrum of dibenzyl disulfide detected by GC of the reaction mixture.



Figure S7. Mass spectrum (GC-MS) of dipentyl disulfide detected by GC of the reaction mixture.



Figure S8. ¹H NMR confirmed the structures of products of disulfides (A) didodecyl disulfide, (B) dibenzothiazole disulfide and (C) dibenzyl disulfide.



Figure S9. Photos of GO (A) and active carbon (B) in round flask. Photos of reaction mixture in round flask with GO (C) and active carbon (D) after deodorization.



Figure S10. GC diagrams of (A) *n*-dedocyl mercaptan solution in heptane, (B) reaction mixture after dedorization with GO and (C) reaction mixture after deodorization with active carbon.



Figure S11. Increase in the ratio of disulfides to mercaptans along with reaction time in heptane. The lines are drawn only for guidance.

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Reaction time (min)	-OH	O-C	O=C	0-C(0)	Quinone
GO	9.5	14.5	2.9	2.2	1.2
120	5.1	9.1	3.1	3.0	1.5
300	3.4	2.2	3.4	2.4	1.1
660	3.3	1.2	2.1	1.4	1.2

Table S1 Fitting results (at.%) of O1s XPS spectra of GO and RGO samples