

Electronic Supplementary Information

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A facile green strategy for rapid reduction of graphene oxide by metallic zinc

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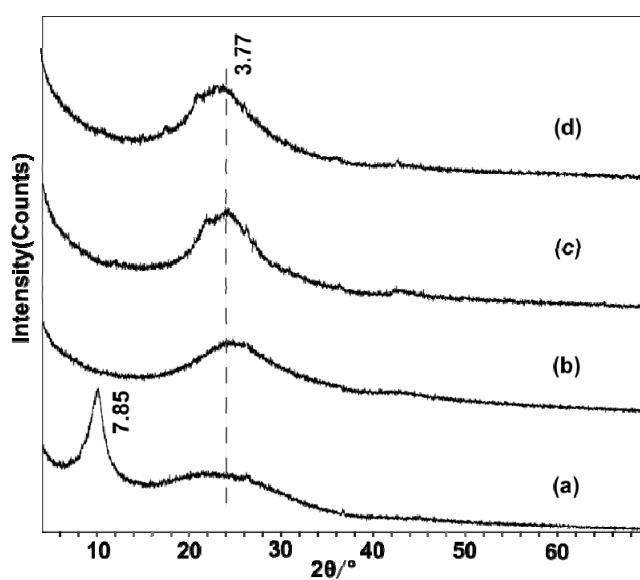


Fig. S1 XRD patterns of GO-NaOH-RT (a), GO-NaOH-100 (b), GO-Zn-RT (c) and GO-Zn-100 (d), respectively. The d -value is in Å.

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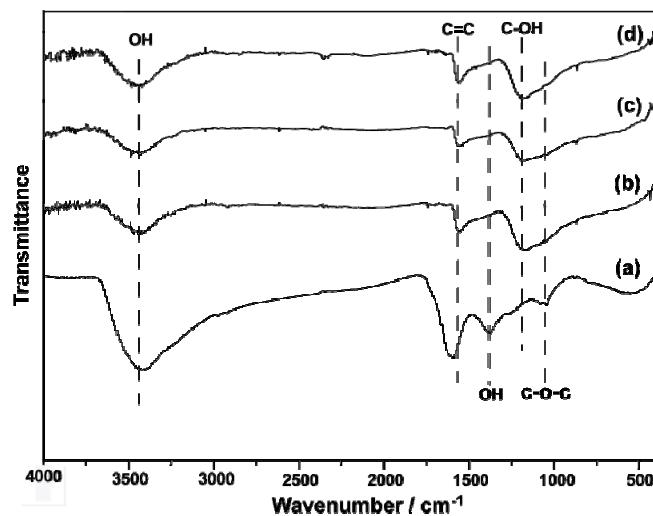


Fig. S2 FT-IR spectra of GO-NaOH-RT (a), GO-NaOH-100 (b), GO-Zn-RT (c) and GO-Zn-100 (d), respectively.

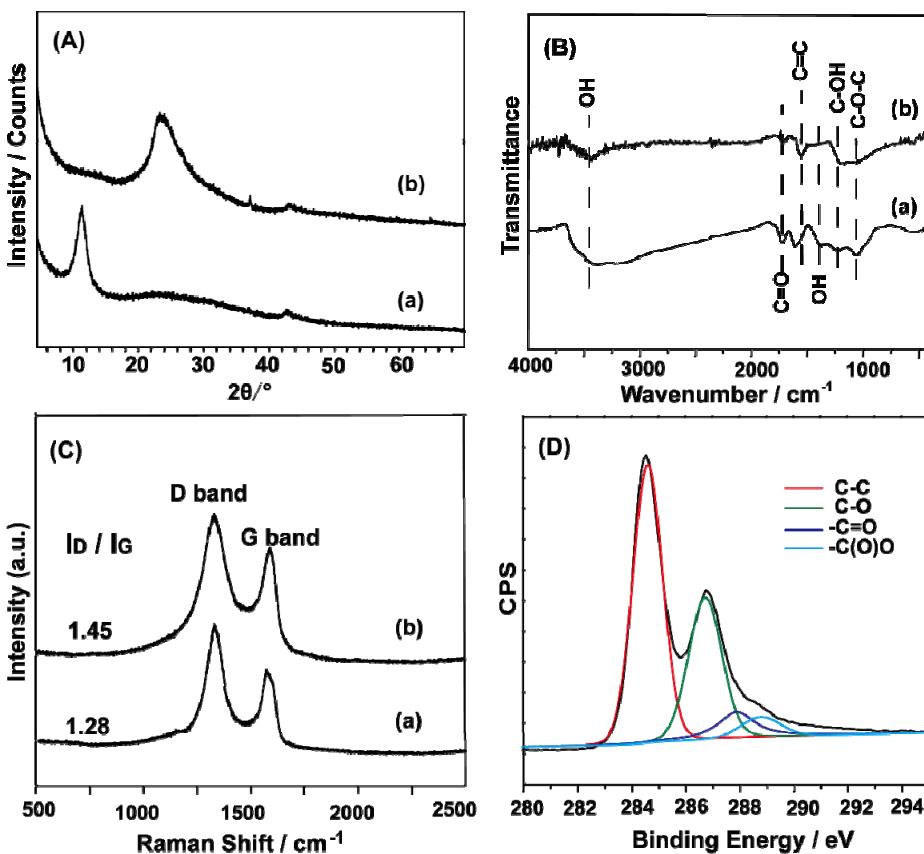


Fig. S3 Control study on the reduction of GO in acidic media (HCl, 1M) at RT and 100 °C, rGO-HCl-RT and rGO-HCl-100 were confirmed by XRD pattern (A), FT-IR (B) and Raman (C) spectra, respectively. (D) is the XPS spectrum of rGO-HCl-100.

We also reduced GO in HCl (1 M) solution at room temperature and 100°C for control experiments, powder of 0.01 M Zn and 0.06 M HCl were added into GO dispersion (0.5 g/L, 60 mL), and the products were donated as rGO-HCl-RT and rGO-HCl-100. A board diffraction peak at about 25° indicates that reduction has taken place at 100 °C and some oxygen-containing groups (ie. carboxyl and epoxy) are removed. According to Raman spectra, rGO-HCl-100 becomes more defective than that of rGO-HCl-RT. However, XPS analysis shows that some oxide groups are still retained and the calculated C/O ratio is 3.06, indicating a low reduction extent of rGO-HCl-100.

This can be clarified as follows. In 1 mol/L of HCl solution, the pH value of the solution is 0.18, well-dispersed GO sheets trend to form agglomeration,¹ They are apt to be captured by the rising hydrogen bubbles and then float to the water surface due to their amphiphilicity,² thus, GO cannot get fully in touch with Zn powder though constant stirring is provided. In addition, the fast reaction rate of Zn powder with HCl acid results in a rapid consumption of Zn. While in 100 °C, as a strong Lewis acid, zinc chloride is known to catalyze the dehydration of alcohols and activate aromatic nucleus.^{3,4} Chemisorbed hydrogen has been noticed to open epoxy and reduce hydroxyl on GO,⁵ that is, generated hydrogen in the solution will also contribute to the reduction.

The mechanism is quite different from that in NaOH solution, it deserves to be noticed.

References

- 1 S. J. Guo and S. J. Dong, *Chem. Soc. Rev.*, 2011, **40**, 2644-2672.
- 2 J. Kim, L. J. Cote, F. Kim, W. Yuan, K. R. Shull and J. X. Huang, *J. Am. Chem. Soc.*, 2010, **132**, 8180-8186.
- 3 J. G. Smith and G. F. Wright, *J. Org. Chem.*, 1952, **17**, 1116-1121.
- 4 B. K. Bettadaiah and P. Srinivas, *Tetrahedron Lett.*, 2003, **44**, 7257-7259.
- 5 S. Kim, S Zhou, Y. K. Hu, M. Acik, Y. J. Chabal, C. Berger, W. Heer, A. Bongiorno and E. Riedo, *Nature Materials*, 2012. doi: 10.1038/NMAT3316