

Electronic Supplementary Information (ESI) for

One-pot scalable synthesis of Cu-CuFe₂O₄/graphene composites as anode material of lithium-ion batteries with enhanced lithium storage properties

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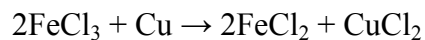
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Preparation of CuFe₂O₄/G composites

A certain amount of Cu-CuFe₂O₄/graphene composites was added into deionized water under sonication to form colloidal suspension, and then FeCl₃ was dissolved into the resultant suspension under mechanical stirring for several hours. The color of the solution became light green due to the presence of FeCl₂ and CuCl₂. The metallic Cu was dissolved by the following reaction:



After washing several times by alcohol and deionized water, the residual powder was collected by centrifugation and dried in oven at 60 °C for 12 h.

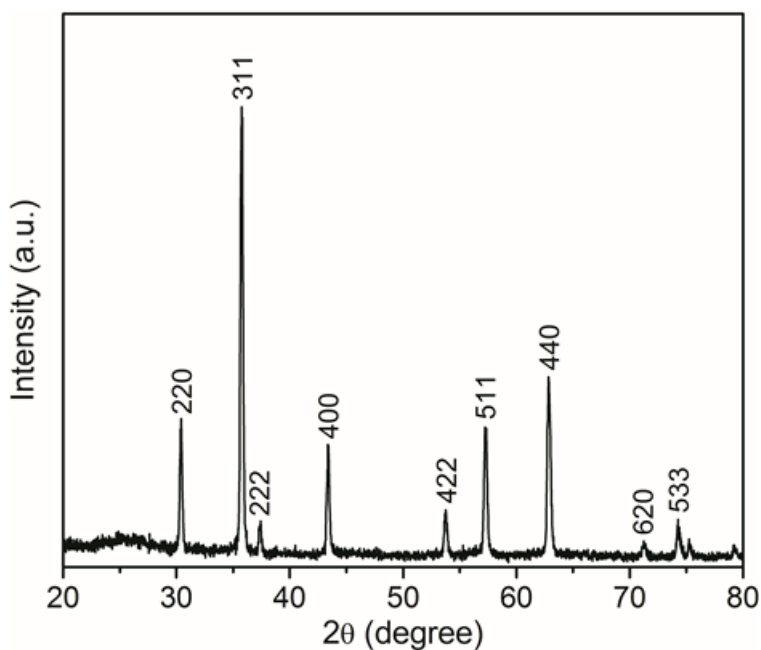


Fig. S1 XRD pattern of FeCl₃ treated Cu-CuFe₂O₄/G composites, all the diffraction peaks and relative intensity are consistent with those cubic phase of CuFe₂O₄ (PDF, 01-077-0010), which confirmed the nonexistence of metallic Cu in the FeCl₃ treated Cu-CuFe₂O₄/G composites and the good crystallinity of CuFe₂O₄/G composites.

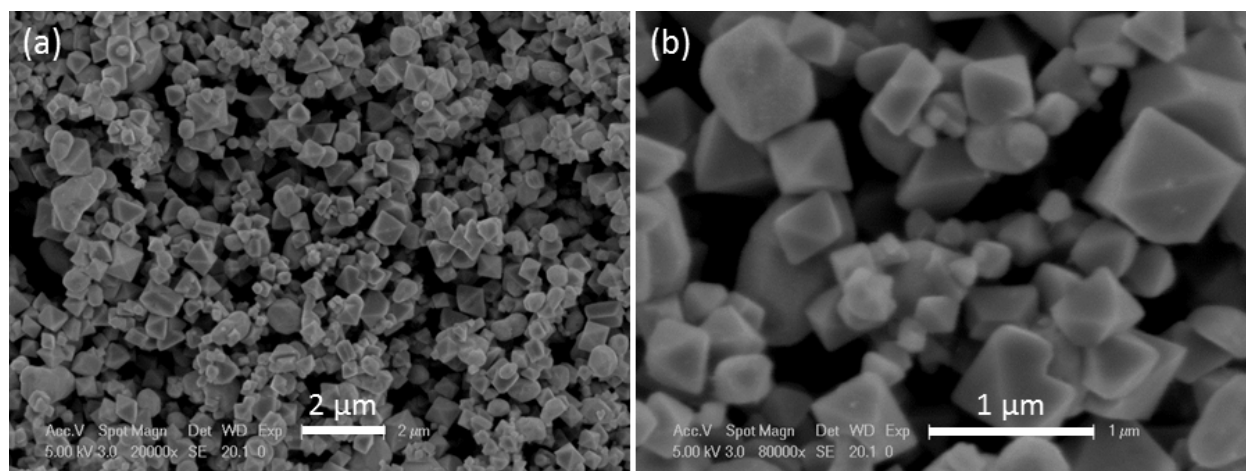


Fig. S2 SEM images of pure Cu-CuFe₂O₄ crystals with different magnification.

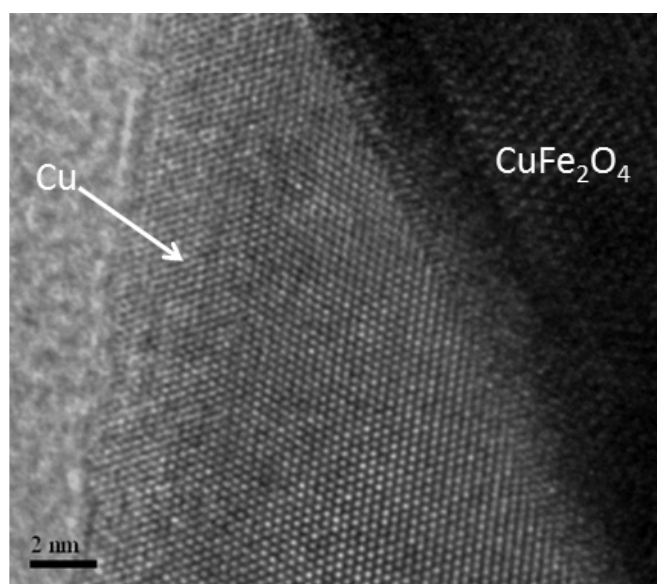


Fig. S3 HRTEM image of metallic Cu attached to the edge of CuFe₂O₄ hexagonal platelet in the Cu-CuFe₂O₄/G composites.

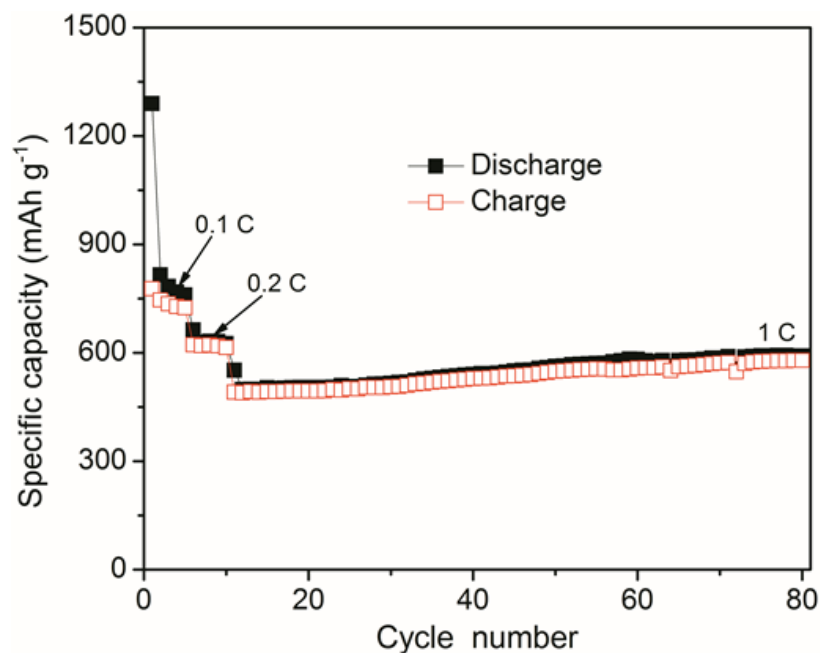


Fig. S4 Cycling performance of CuFe₂O₄/G composites at a current density of 1000 mA/g. (1C=1000 mA/g)

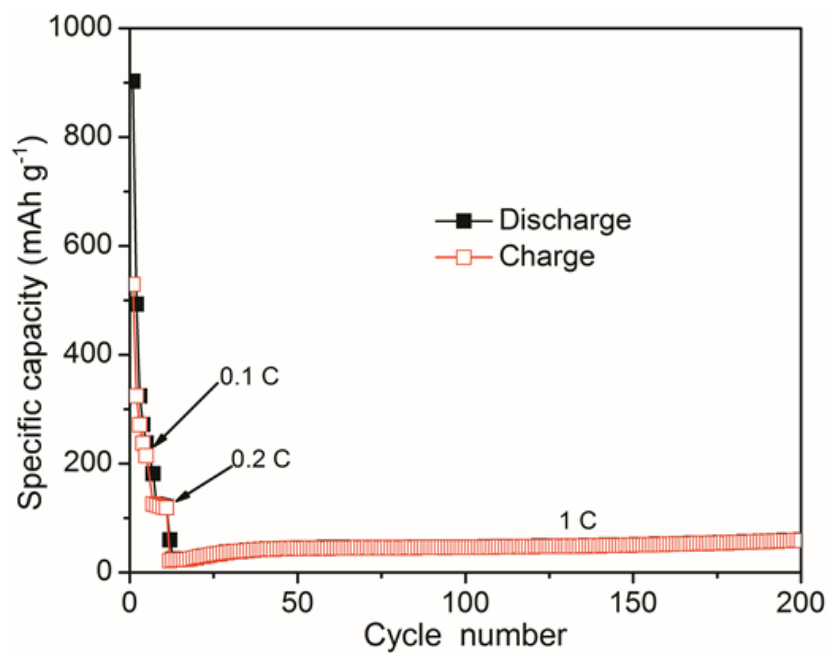


Fig. S5 Cyclic stability of pure Cu-CuFe₂O₄ crystals at a current density of 1000 mA/g. (1C=1000 mA/g)