People's Republic of China.

## Rational design of metal organic frameworks-derived FeS<sub>2</sub> hollow nanocages@reduced graphene oxide for K-ion storage

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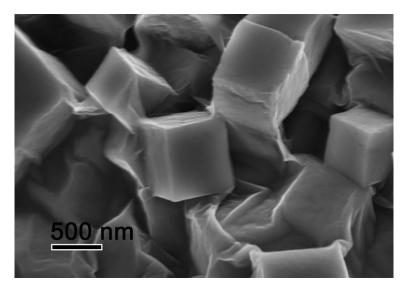


Figure S1 SEM image of PB@GO.

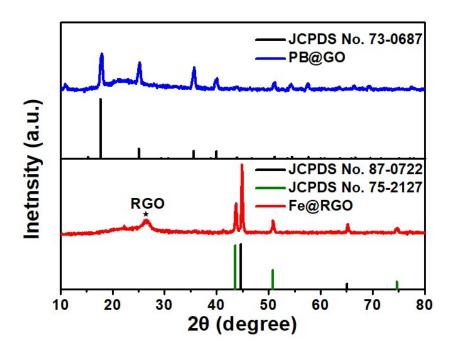
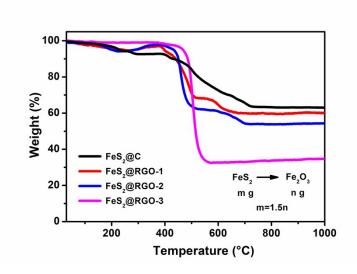


Figure S2 XRD patterns of PB@GO and Fe@RGO.

In the XRD pattern of Fe@RGO, a broad peak at 26° is attributed to (002) facets of RGO. Other diffraction peaks are assigned to Fe (JCPDS card No. 87-0722) and FeN<sub>0.0324</sub> (JCPDS card No. 75-2127). The result demonstrates that the RGO and Fe have been reduced after thermal treatment. The formation of FeN<sub>0.0324</sub> is mainly due to the existence of N element in PB. According to the previous reports, during annealing progress in N<sub>2</sub> or Ar atmosphere, transition metal anions will be reduced into pure transition metal, and CN- group linkers will form carbon compounds [1-3]. Furthermore, RGO is also an excellent reducing agent in high temperature, which is helpful for the reduction of Fe ion. Such phenomenon has been reported in other Co-Fe alloys, Ni-Fe alloys [1-3].



**Figure S3** TGA curves of FeS<sub>2</sub>@C, FeS<sub>2</sub>@RGO-1, FeS<sub>2</sub>@RGO-2 and FeS<sub>2</sub>@RGO-3.

TGA curves were carried out to evaluate the carbon content in the composite, as shown in Fig. S3. After being heated to 700 °C, the weights of the samples become stable, and 63.4%, 59.0%, 53.8% and 32.9% of the original weight are left for the final products for FeS<sub>2</sub>@C, FeS<sub>2</sub>@RGO-1, FeS<sub>2</sub>@RGO-2, FeS<sub>2</sub>@RGO-3, respectively. Due to the oxidation of FeS<sub>2</sub> to Fe<sub>2</sub>O<sub>3</sub> while the carbon to carbon dioxide, the carbon content of FeS<sub>2</sub>@C, FeS<sub>2</sub>@RGO-1, FeS<sub>2</sub>@RGO-2, FeS<sub>2</sub>@RGO-3 can be calculated, and corresponding values are 4.9%, 11.5%, 19.3% and 50.7%.

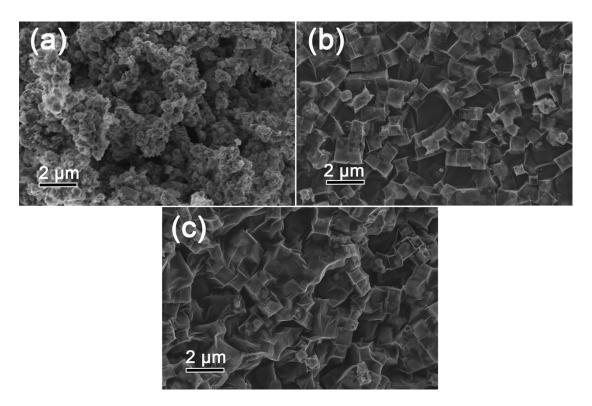
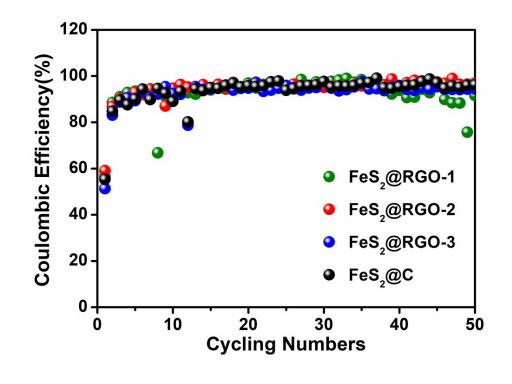


Figure S4 SEM images of (a) FeS<sub>2</sub>@C, (b) FeS<sub>2</sub>@RGO-1 and (c) FeS<sub>2</sub>@RGO-3



**Figure. S5** Coulombic efficiencies of FeS<sub>2</sub>@C, FeS<sub>2</sub>@RGO-1, FeS<sub>2</sub>@RGO-2 and FeS<sub>2</sub>@RGO-3 at 50 mA  $g^{-1}$ .

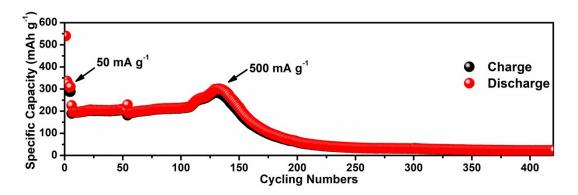


Figure S6 Long-term cycling performance of FeS<sub>2</sub>@RGO-1 at 500 mA g<sup>-1</sup>

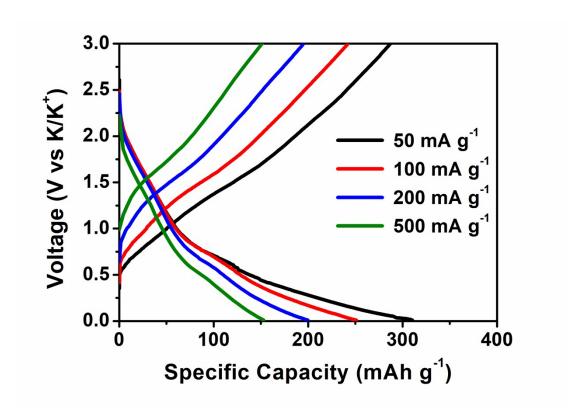
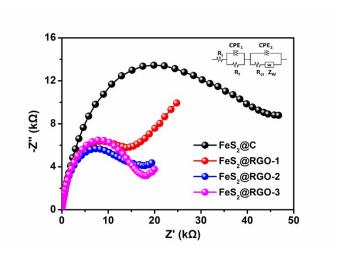
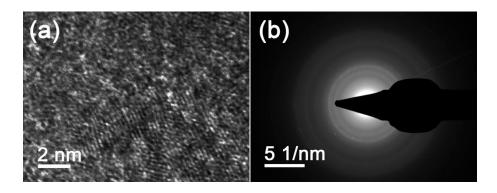


Figure S7 Galvanostatic charge-discharge curves of  $FeS_2@RGO-2$  electrode at different current densities.



**Figure S8** EIS of FeS<sub>2</sub>@C, FeS<sub>2</sub>@RGO-1, FeS<sub>2</sub>@RGO-2 and FeS<sub>2</sub>@RGO-3 after 10 cycles.



**Figure S9** (a) HRTEM image of FeS<sub>2</sub>@RGO-2 electrode when charging to 3.0 V and (b) the corresponding SAED image in the first cycle.

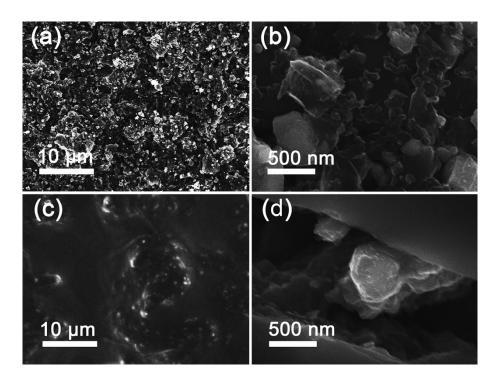


Figure S10 SEM images of  $FeS_2@RGO-2$  electrodes. (a)-(b) before cycles, (c)-(d) after a number of cycles.

## Reference

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