Supporting information

Facile synthesis and catalytic performance of Co₃O₄ nanosheets

in-situ formed on reduced graphene oxide modified Ni foam

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a) b) Co,O/rGO GO Co₃O₄/rGO 185 662 Intensity (a.u.) 490 Intensity (a.u.) 500 1000 1500 2000 2500 100 200 300 400 500 600 700 Wave number (cm⁻¹) Wave number (cm⁻¹)

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Figure S1 Raman spectra of Co_3O_4/rGO and GO(a) and partial enlargement(b) of Figure S1(a)

Raman spectra of Co_3O_4/rGO and GO were measured and shown in the following Figure S1. On both spectra of Co_3O_4/rGO and GO, two remarkable peaks located at 1349 and 1598 cm⁻¹ referring to the D and G bands respectively. The D band is arose from the edge or defect sites of carbon and the G band corresponds to ordered sp²bonded carbon atoms. As seen from Figure S1(a), the spectra of Co_3O_4/rGO and GO are very similar. When enlarging the region in Figure S1(a) marked by green rectangle, it can be found from Figure S1(b) that three peaks at 185, 490 and 670 cm⁻¹ which can be assigned to the Eg, F_2g and A_1g modes of Co_3O_4 indicating that $Co_3O_4/rGO@Ni$ foam electrode was successfully prepared.



Figure S2. The N_2 adsorption/desorption isotherms of $Co_3O_4(a)$ and $Co_3O_4/rGO(b)$.

To investigate the surface area of Co_3O_4 @Ni foam and Co_3O_4 /rGO@Ni foam, N₂ isothermal adsorption-desorption measurements were conducted. The samples of Co_3O_4 and Co_3O_4 /rGO were obtained by scraping the powders from the Ni foam substrate. As seen from Figure S2, the N₂ adsorption-desorption isotherms of both Co_3O_4 and Co_3O_4 /rGO exhibit typical IV isotherms, indicating the mesoporous structures of Co_3O_4 and Co_3O_4 /rGO. In addition, the results show a relatively high specific surface area of 55.45 m² g⁻¹ of Co_3O_4 /rGO which is higher than that of Co_3O_4 (48.06 m² g⁻¹). Thus, it can be concluded that the addition of rGO efficiently increases the surface area of the electrode and then leads to a better catalytic activity of the Co_3O_4 /rGO@Ni foam electrode than Co_3O_4 @Ni foam electrode.