# **High performances aqueous rechargeable nickel//bismuth batteries with Bi2MoO6@rGO and Co0.5Ni0.5MoO4@rGO as electrode materials**

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## **Electrochemical Calculations:**

## **1.Single electrode:**

Based on discharge curves, the specific gravimetric capacity of active materials for the single electrode can be calculated according to the following equation (1):

$$
C_m = \frac{I}{m} \times \Delta t \tag{1}
$$

Where  $C_m$  (mAh/g) means the specific gravimetric capacity,  $I(A)$  means the discharge current,  $m$  (g) means the mass of active materials in the single electrode,  $\Delta t$ (s) means discharge time.

### **2. Ni//Bi battery:**

The mass of the electrode active materials should be adjusted before assembling Ni//Bi battery so that two electrodes are balanced on the basis of equation (2):

$$
C_m^- \times M_{-} = C_m^+ \times M_{+}
$$

(2)

Where  $C_m$  and  $C_m$ <sup>+</sup> are the specific capacity of the cathode and anode (mAh/g),  $M$  + and  $M$  - are the mass of active materials (mg), respectively.

The specific capacity of two-electrode cell model  $(C_{cell})$  can also be calculated depending on software or equation (3):

$$
C_{cell} = \frac{I}{m} \times \Delta t
$$
\n(3)

Where  $C_{Cell}$  (mAh/g) is specific capacity, *I* (A) is discharge current,  $\Delta t$  (s) is discharging time, and m (g) is total mass of active materials.

The energy density (*E*) and power density (*P*) of Ni//Bi battery can be calculated by the following equations (4) and (5):

$$
E = \int_{0}^{\Delta t} IV_{(t)} dt
$$
  
\n
$$
P = \frac{E}{\Delta t}
$$
 (4)

Where  $E$  (Wh/kg) is energy density,  $I(A/g)$  is current density,  $V(t)$  (V) is discharging voltage and  $dt$  is time differential,  $P$  (W/kg) is power density,  $\Delta t$  (s) is the discharging time.

### **The calculation of "b values":**

The relationship between peak current (i) and scanning rate (v) obeying the equation (6):

$$
i=av^b\tag{6}
$$

Where a and b are different positive number. When  $0.5 < b < 1$ , the electrochemical behavior of electrode active materials is controlled by diffusion-controlled process and surface-capacitive effect. When b=0.5, the electrochemical behavior is controlled by diffusion process. And when b=1, the electrode exhibits surface-capacitive effect.



**Fig.** S1 CV curves of  $Bi_2MoO_6@rGO-25$  (a); The relation between the anodic/cathodic peak current and the scan rate of  $Bi<sub>2</sub>MoO<sub>6</sub>(a)<sub>r</sub>GO-25$  (b)



**Fig.** S2 CV curves of  $Bi_2MoO_6@rGO-10$  (a) and  $Bi_2MoO_6@rGO-50$  (c); GCD curves of  $Bi<sub>2</sub>MoO<sub>6</sub>(QrGO-10 (b) and Bi<sub>2</sub>MoO<sub>6</sub>(QrGO-50 (d))$ 



**Fig.** S3 XPS survey spectrum of  $Co<sub>0.5</sub>Ni<sub>0.5</sub>MoO<sub>4</sub>@rGO-10$ 



**Fig. S**4 SEM image of graphene oxide for cathode active materials



**Fig. S**5 The relation between the anodic/cathodic peak current and the scan rate of  $Co<sub>0.5</sub>Ni<sub>0.5</sub>MoO<sub>4</sub>@rGO-10$  (a); Cycling performance of  $Co<sub>0.5</sub>Ni<sub>0.5</sub>MoO<sub>4</sub>@rGO-10$  at 15 A/g (b)



**Fig.** S6 CV curves of  $Co_{0.5}Ni_{0.5}MoO_4@rGO-5$  (a) and  $Co_{0.5}Ni_{0.5}MoO_4@rGO-15$  (c); GCD curves of  $Co_{0.5}Ni_{0.5}MoO_{4}@rGO-5$  (b) and  $Co_{0.5}Ni_{0.5}MoO_{4}@rGO-15$  (d)



**Fig. S**7 Discharging curves at different current densities (a); picture shows two devices connected in series can light up one red LED bulb (b)