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ARTICLE

Title: Rechargeable Li//Br battery: A promising platform for post lithium ion batteries Authors: Zheng Chang, Xujiong Wang, Yaqiong Yang, Jie Gao, Lili Liu, Minxia Li, Yuping Wu\*

# **Electronic supporting information (ESI)**



1. Ionic conductivity of aqueous positive electrode (LiBr aqueous solution added with Br<sub>2</sub>)

Figure S1. (a) Ionic conductivity of LiBr aqueous solution of different concentration, (b) and (c) ionic conductivities of 1 and 7 mol  $L^{-1}$  LiBr aqueous solutions adding with different concentration of Br<sub>2</sub>.

The ionic conductivity at room temperature increases with the concentration of LiBr, and arrives at the highest, 197.3 mS cm<sup>-1</sup>, for 7.76 mol L<sup>-1</sup>.

After adding  $Br_2$  in the LiBr aqueous solution, the ionic conductivity decreases with the concentration of  $Br_2$  due to the transformation of  $Br^-$  into  $Br^{3-}$  or  $Br^{5-}$ .

## 2. Changes of LiBr solution acting as electrolyte before and after charge



Figure S2. (a) Optical images of the solution around the GC electrode surface before and after a charge in 3 mol  $L^{-1}$  LiBr aqueous solution, and (b) UV-Vis spectra of the solution before (black) and after (red) charge.

In Fig.S2a, it was found that the color of the solution after a charge was turned into yellowish, proving the formation of  $Br_3$ . After the charge, the characteristic absorption peak at 274 nm in the UV-Vis spectra was observed, also evidencing the generation of  $Br_3$ .

#### 3. Electrochemical window of GC electrode



Figure S3. CV curves of the GC electrode in the aqueous solutions at the scan rate of 1 mV s<sup>-1</sup>.

It suggests that the electrochemical window in 0.5 mol  $L^{-1}$  Li<sub>2</sub>SO<sub>4</sub> (black) is very wide, and there are no evident redox reactions between -1.5 and 1.5 V (vs. SCE). In the case of 7 mol  $L^{-1}$  LiBr, there is no oxygen evolution, and only the redox for Br/Br<sub>3</sub><sup>-</sup> can be observed.

### 4. Stability of Br electrode



Figure S4. CV curves of 7mol L<sup>-1</sup> LiBr aqueous solution at the scan rate of 0.5 mV s<sup>-1</sup> for the first 10 scans.

The good overlapping of the CV curves demonstrates the excellent stability of Br electrode during the electrochemical reaction.

#### 5. A promising platform of charge and discharge



Figure S5. The charge /discharge curves for 100% DOD of the Li//Br battery in which aqueous positive electrode is 1 mol  $L^{-1}$  LiBr with 0.1 mol  $L^{-1}$  Br<sub>2</sub>.

#### 6. Discharge voltage curves in different current density



Figure S5. Discharge voltage curves in the current density range of 0.42 to 12.7 mA cm<sup>-2</sup>. At each current density, the test was carried out for 5 mins instead of several seconds.



Figure S6. Discharge curves for Br single electrode (1 mol L<sup>-1</sup> Br<sub>2</sub>) in 7 mol L<sup>-1</sup> LiBr at different current densities.

When the current density increases from 0.5 to 5 mA cm<sup>-2</sup>, the voltage decrease is only about 0.3 V, indicating excellent electrochemical redox kinetics.

## 7. The stability of both the solid state electrolyte film



Figure S7. SEM micrographs of LISICON layer before and after 100 cycles at 35% depth of discharge.

# 8. Comparison between different aqueous rechargeable lithium batteries

System	Current density / mA cm <sup>-2</sup>	Average discharge voltage / V	Highest power density / mW cm <sup>-2</sup>	Reference*
Li//I	12	2.9	34.8	16
Li//Fe(CN) <sub>6</sub> <sup>3-</sup>	6.5	1.8	12.5	22
Li//Br	12.3	2.5	29.7	This work

Table S1. Some data about the highest power densities for different aqueous rechargeable lithium batteries.

\*: In the list of references of the main context.