

Supporting Information

**Salt sealing strategy to prepare N, O-codoped
porous bio-carbon derived from Ephedra Herb for
supercapacitor**

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Electrochemical Measurements

1. Fabrication of three-electrode system

In a three-electrode system, the tested sample was loaded onto a normal the glassy carbon electrode (5 mm diameter). Generally, accurately weighed (4.0 mg) sample was ultrasonically dispersed in 0.400 mL Nafion solution (0.25%, DuPont, USA). 8.0 μL of above suspension was dropped onto the working electrode surface and dried at room-temperature (about 0.4 mg cm^{-2}). All of the electrodes were immersed in 6 mol L^{-1} KOH electrolyte. The electrochemical performances were investigated by cyclic voltammetry (CV) and galvanostatic charge/discharge measurements from CHI 660E electrochemical workstation. The charge-discharge cycle stability was employed on a LAND CT2001A (Wuhan Land Instrument Company, China).

2. Fabrication of two-electrode supercapacitor

During the two-electrodes test, the electrodes were made by mixing S_{12} (80 wt%), polyvinylidene fluoride (PVDF, 10 wt%) and commercial carbon black (10 wt%). The mixture was coated on nickel foam and then pressed at 15 MPa followed by drying at 393 K for 12 h. A button-type supercapacitor was assembled with two similar carbon electrodes (about 2.6 mg cm^{-2}) separated by a polypropylene membrane in 6 M KOH electrolyte and 1 M Li_2SO_4 . The charge-discharge performance was measured using a CHI 760E electrochemical workstation (CH Instrument, Shanghai, China). The specific capacitance of the S_{122} material (C, in F g^{-1}) was calculated based on the discharge curve according to equation (1).

$$C = \frac{2I}{m \frac{\Delta V}{\Delta t}} \quad (1)$$

Where I is the discharge current (A), $\frac{\Delta V}{\Delta t}$ ($V s^{-1}$) is the slope obtained by fitting a straight line to the discharge voltage, and m is the mass (g) of active material in a single electrode.

The energy density (E , in $W h kg^{-1}$) and average power density (P , in $W kg^{-1}$) were calculated according to equation (2) and (3).

$$E = \frac{1}{2 * 4 * 3.6} C V^2 \quad (2)$$

$$P = \frac{E}{\Delta t d} \quad (3)$$

Where C is the specific capacitance of a single electrode in a two electrode supercapacitor ($F g^{-1}$), and V is the usable voltage after IR drop (V), and $\Delta t d$ is the discharge time (h). [S1-S5]

3. Materials Characterization

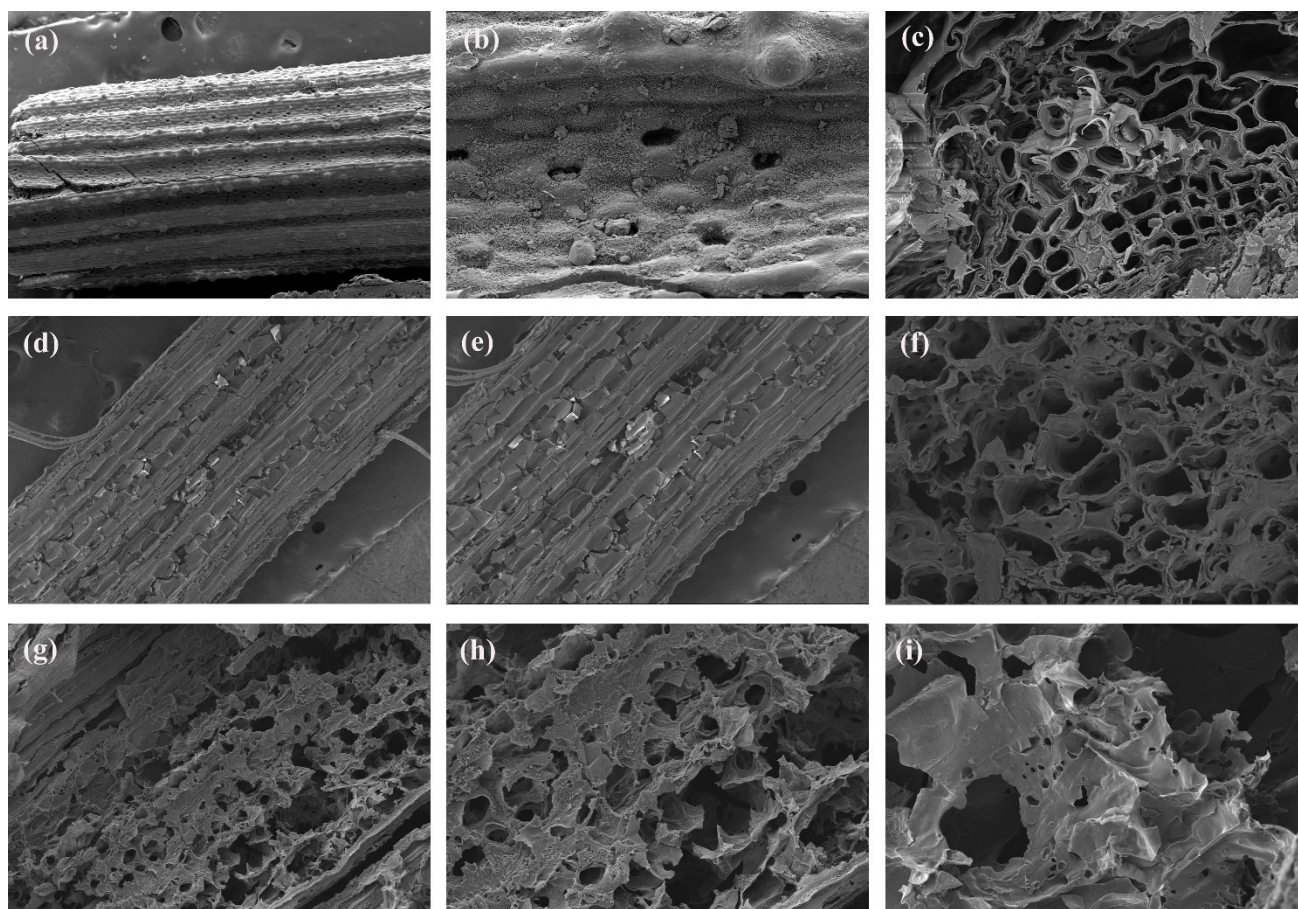


Fig. S1 SEM and HR-SEM images of the S₁₀ (a, b, c), S₁₁ (d, e, f) and S₁₂ (g, h, i).

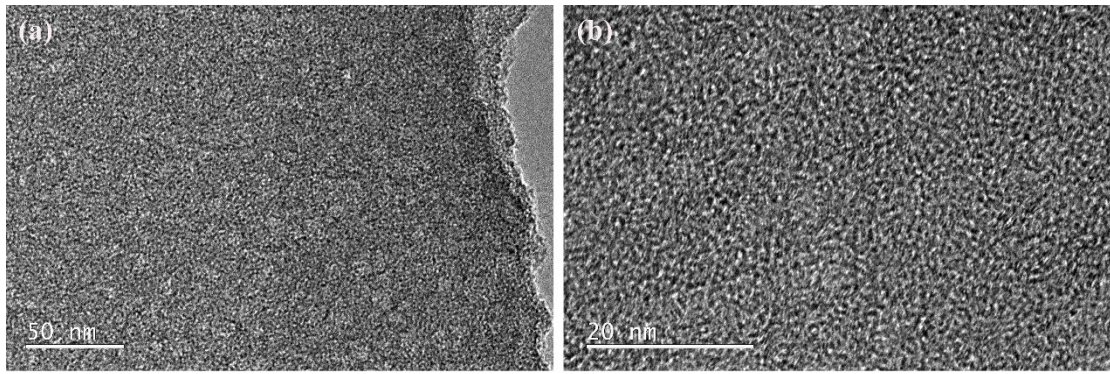


Fig. S2 TEM and HR-TEM images of the S₁₂ (a, b).

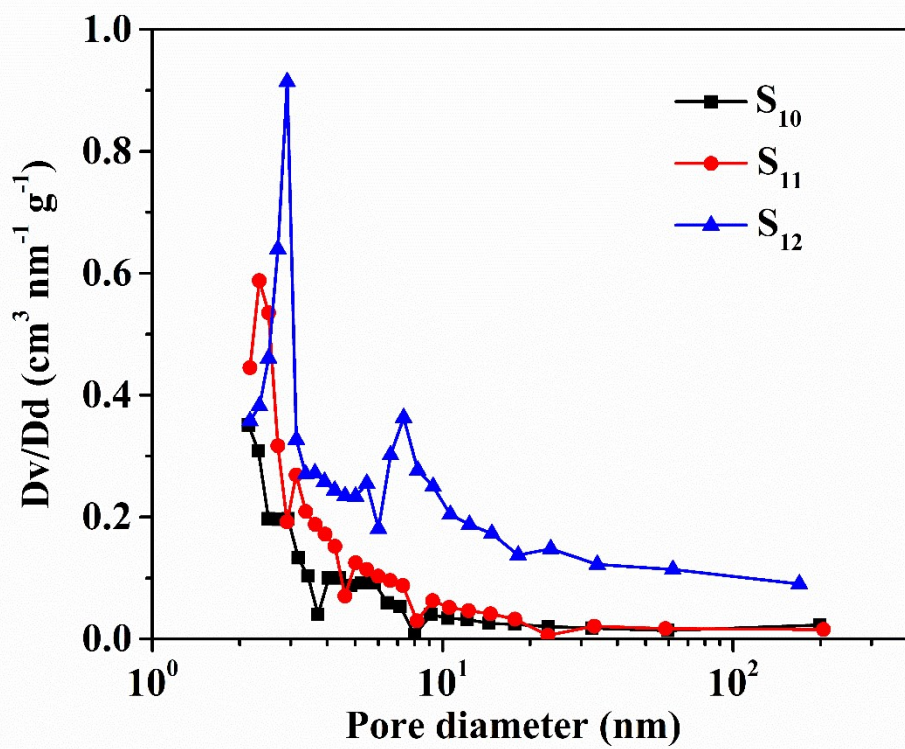


Fig. S3 Pore size distribution of S_{10} , S_{11} and S_{12} .

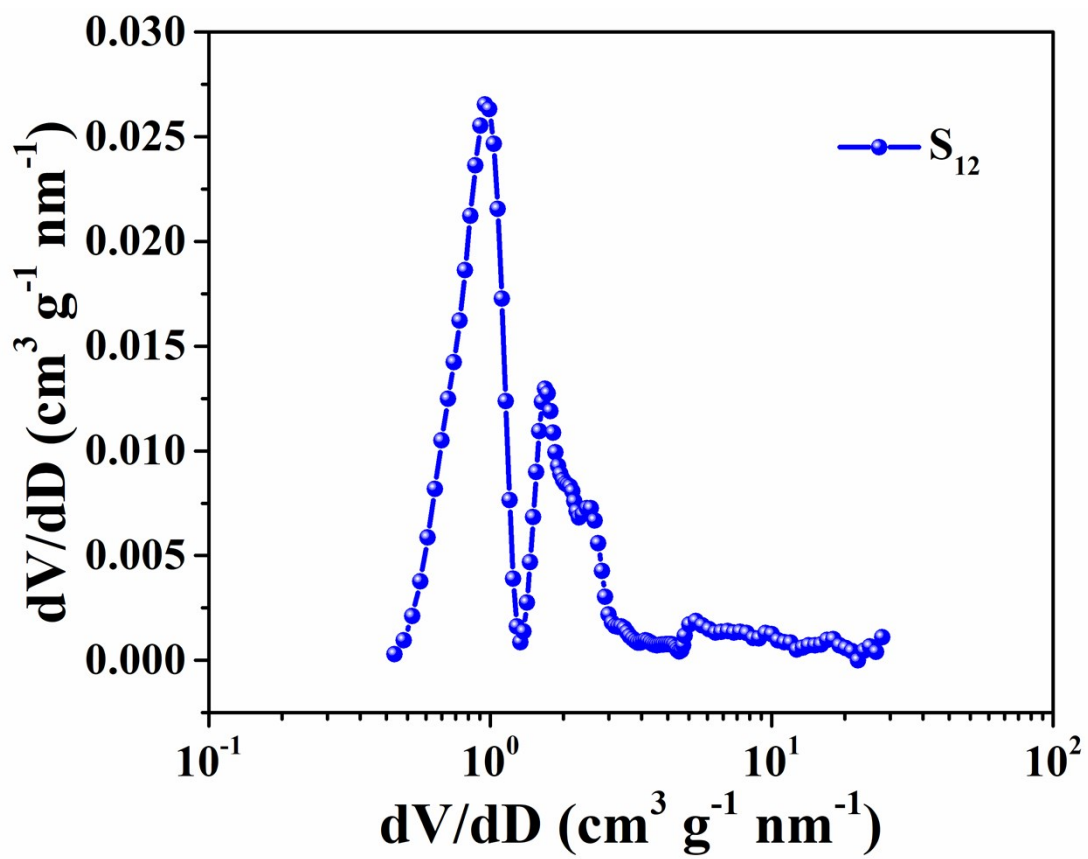


Fig. S4 The S_{12} samples was further verified by the density functional theory (DFT) method.

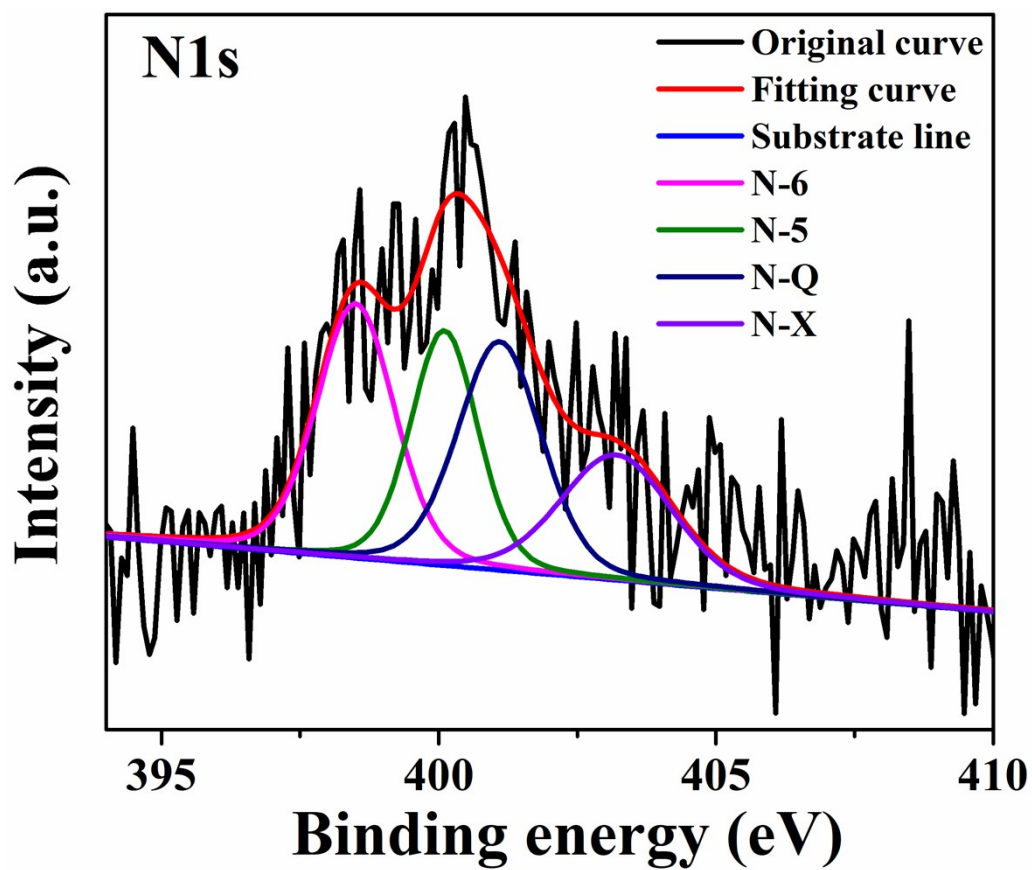


Fig. S5 High-resolution X-ray photoelectron spectroscopy (XPS) of N1s region for S₁₂ sample.

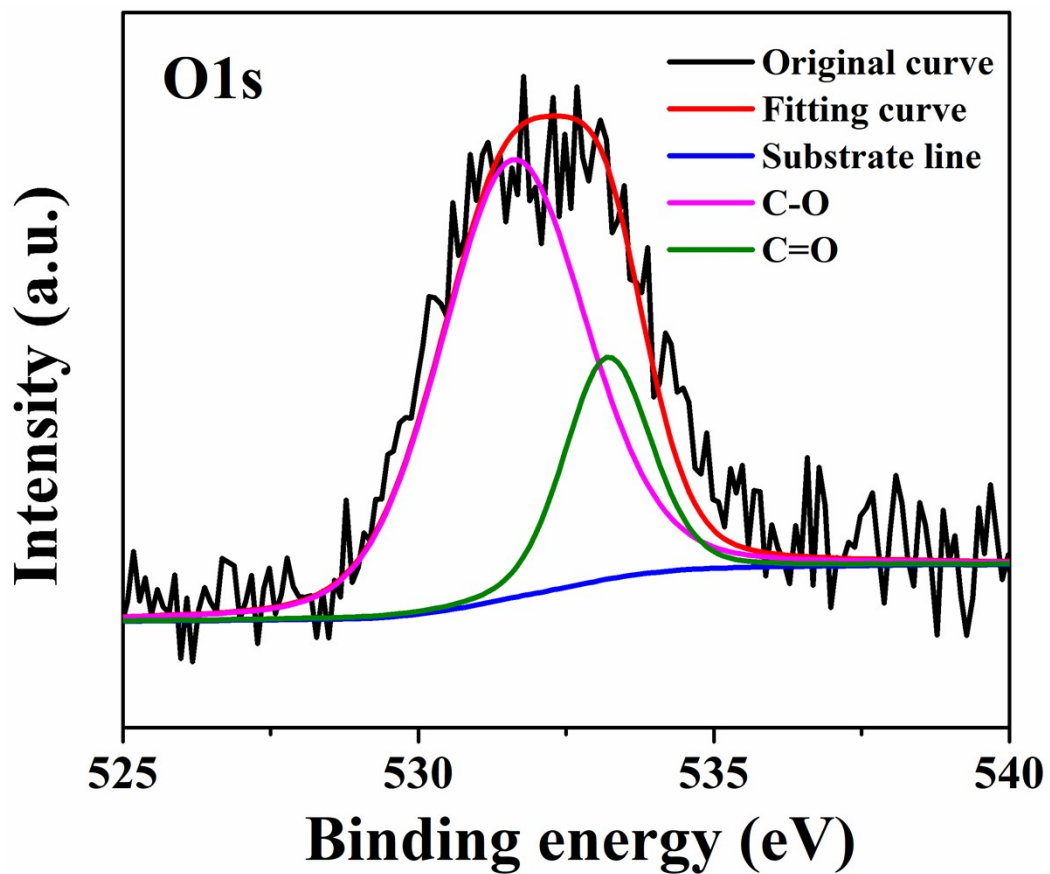


Fig. S6 High-resolution X-ray photoelectron spectroscopy (XPS) of O1s region for S₁₂ sample.

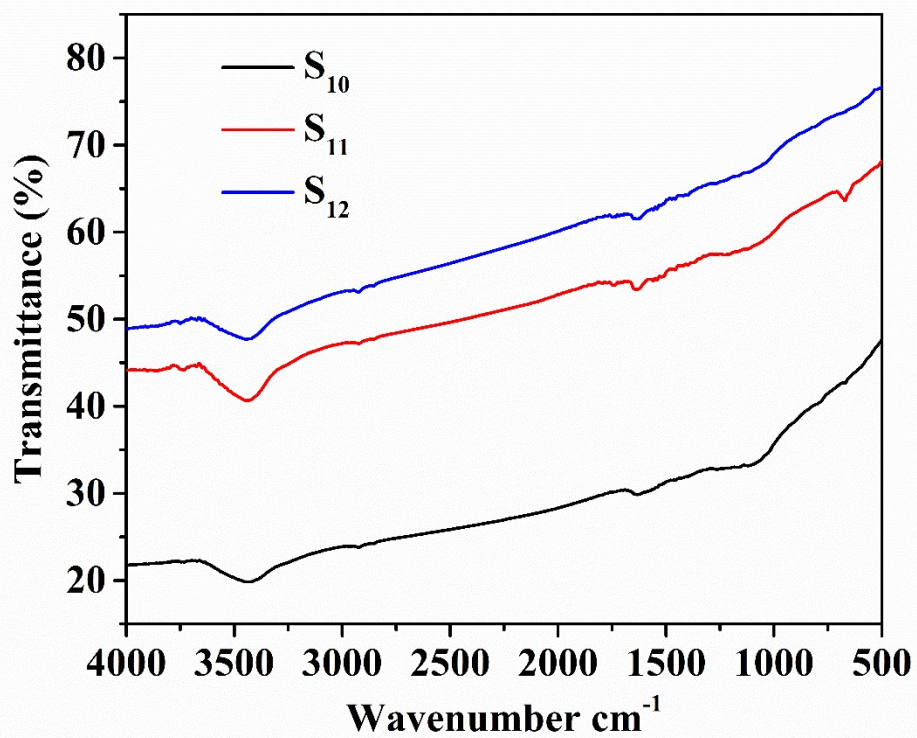


Fig. S7 FT-IR analysis of the obtained materials.[S6, S7]

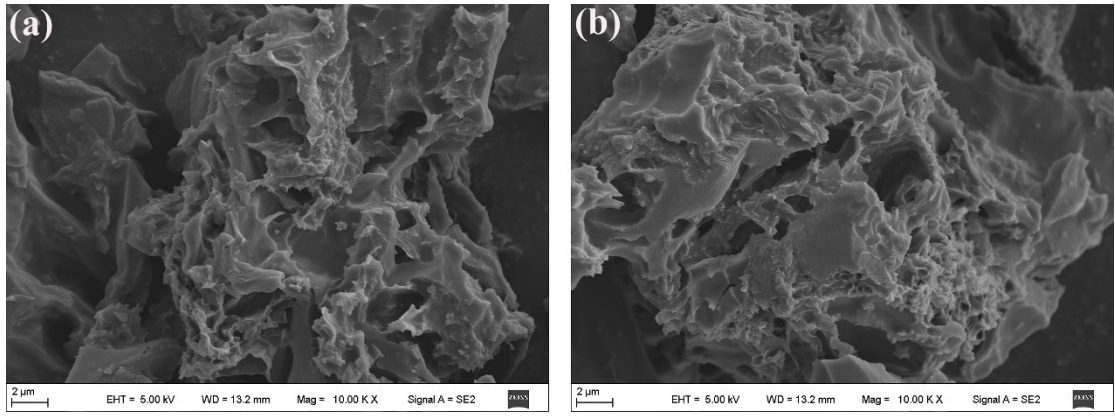


Fig. S8 The SEM of S₁₂ after cycling.

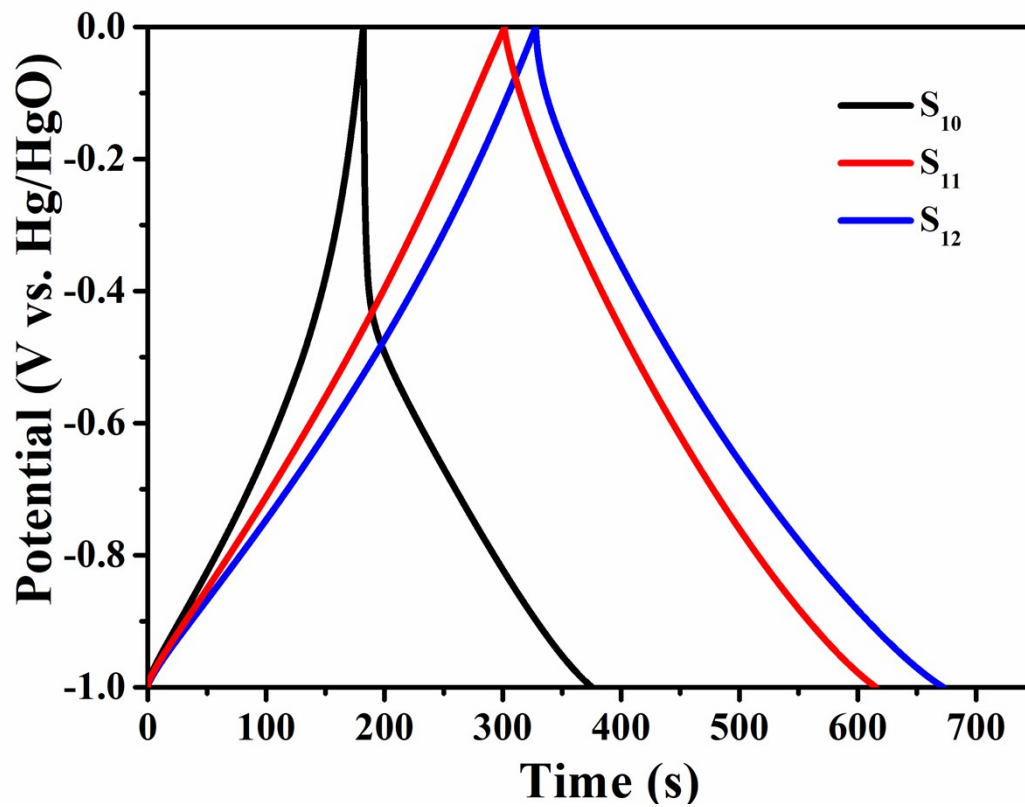


Fig. S9 GCDs patterns of S₁₀, S₁₁, S₁₂ electrode at current density of 0.5 A g⁻¹.

References

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