

Supporting Information

Conductive 1D High-nucleus Silver polymer as a Brilliant Non-hybrid Supercapacitor Electrode

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1. Electrochemical measurements

The cyclic voltammetry (CV) measurement like other electrochemical characterization includes of galvanostatic charge-discharge (GCD) and electrochemical impedance spectroscopy (EIS) was achieved with the same equipment, PalmSens3 electrochemical workstation using a three electrode system at ambient temperature. In which, Ag/AgCl and high-surface-area Pt electrode applied as the reference and counter electrodes, respectively.

The cyclic voltammetry curves were plotted with different scan rates from 200-1000 mVs⁻¹ in potential window between -0.5 to +0.5 V. The EIS spectroscopy were carried out in the frequency range from 50 kHz to 50 mHz in OCP (AC perturbation was 10 mV). The GCD test

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was fulfilled at various current densities (5.0 to 20.0 A/g). The capacities of the electrode materials were calculated by both CV curves and GCD curves (discharge part) based on equation 1 and 2, where ΔQ , ΔV and m are the integrate of voltammetric charges from CV curves, the potential range in the CV (V) and the mass of the sponge electrode (g) for **equation 1** and I , Δt , ΔV and m are the discharge current and time (s), the potential window (V) and mass of the electroactive material (g), respectively in **equation 2**:

$$C_{sp} = \Delta Q / \Delta V \times m \quad (\text{equation 1})$$

$$C_{sp} = I \times \Delta t / \Delta V \times m \quad (\text{equation 2})$$

2. Electrode and device Preparation

The working electrode assembled by mixing the complex and binder (polytetrafluoroethylene) with 95: 5 mass ratios. The obtained viscous slurry after 5 minutes ultrasonication was deposited on a $1 \times 1 \text{ cm}^2$ size Ti foam, then pressed under 10 MPa pressure and dried under vacuum for 10 h at 80 °C before any electrochemical test. It should be mentioned that the weight of active electrode material was in a range of 1-1.5 mg. In order to practical application study, as-prepared SSc electrodes were used as the positive and the negative electrodes in two-electrode cell configuration. A paper soaked and saturated in H_2SO_4 (1M) was placed between two electrodes and both electrodes were completely attached to the paper by clamps. The electrodes were connected to a green LED by two lizard clamp cable. The designed device was used, illustrating that the green LED hold on for few minutes.

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Table S1. Crystal data and structure refinement for **CP₁**

Chemical formula	C ₃₁ H ₃₀ Ag ₈ F ₁₂ NO ₉
formula mass	1651.52 g/mol
<i>T</i> (K)	150(2)
Wavelength	0.71073 Å
Crystal system	Monoclinic
space group	P2(1)/c
<i>a</i> (Å)	15.0304(14)
<i>b</i> (Å)	22.8131(18)
<i>c</i> (Å)	13.6252(13)
α (deg)	90
β (deg)	107.640(3)
γ (deg)	90
V (Å ³)	4452.3(7)
Z	4
$\mu_{\text{calcd.}}$ (g/m ³)	2.464
Absorption coefficient	3.538 mm ⁻¹
F(000)	3116
Reflections collected	61813 / 10222 [R(int) = 0.0421]
Refinement method	Full-matrix least squares on F ²
Goodness-of-fit on F ²	1.032
R indices (all data)	R ₁ ^a = 0.0582, wR ₂ ^b = 0.0923

^a $R_1 = [\sum \text{abs}(\text{abs}(\text{Fo}) - \text{abs}(\text{Fc}))] / [\sum \text{abs}(\text{Fo})]$. ^b $wR_2 = [\sum (w(\text{Fo}^2 - \text{Fc}^2)^2) / \sum [w(\text{Fo}^2)^2]]^{0.5}$.

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Table S5. Selected bond lengths [\AA] and angles [$^\circ$] for CP_1 .

Ag(1)-C(1)	2.117(6)	Ag(4)-O(2)	2.329(4)
Ag(1)-C(21)	2.146(5)	Ag(4)-Ag(2)#2	3.0022(6)
Ag(1)-Ag(4)	2.8360(6)	Ag(5)-N(1)	2.261(6)
Ag(1)-Ag(6)	2.9119(6)	Ag(5)-C(11)#2	2.329(5)
Ag(1)-Ag(8)	2.9679(7)	Ag(5)-O(32)	2.343(5)
Ag(1)-Ag(3)	2.9980(7)	Ag(5)-C(1)	2.587(6)
Ag(1)-Ag(5)	3.1166(7)	Ag(5)-Ag(7)	2.8737(7)
Ag(1)-Ag(7)	3.1744(7)	Ag(5)-Ag(2)#2	3.0794(7)
Ag(1)-Ag(2)	3.2033(6)	Ag(6)-O(22)	2.296(4)
Ag(2)-C(11)	2.149(5)	Ag(6)-C(11)	2.346(5)
Ag(2)-C(31)	2.154(6)	Ag(6)-C(21)	2.409(5)
Ag(2)-Ag(7)#1	2.8962(7)	Ag(6)-C(22)	2.464(6)
Ag(2)-Ag(8)	2.9166(8)	Ag(6)-C(12)	2.642(6)
Ag(2)-Ag(4)#1	3.0022(6)	Ag(6)-Ag(7)#1	3.0051(7)
Ag(2)-Ag(6)	3.0439(7)	Ag(7)-O(14)	2.29(4)
Ag(2)-Ag(5)#1	3.0794(7)	Ag(3)-Ag(1)-Ag(7)	109.646(19)
Ag(2)-Ag(3)	3.1716(7)	Ag(5)-Ag(1)-Ag(7)	54.353(14)
Ag(3)-O(1)	2.277(4)	C(1)-Ag(1)-Ag(2)	77.24(15)
Ag(3)-C(31)	2.314(5)	C(21)-Ag(1)-Ag(2)	95.71(14)
Ag(3)-C(21)	2.403(6)	Ag(4)-Ag(1)-Ag(2)	127.217(19)
Ag(3)-C(32)	2.649(6)	Ag(6)-Ag(1)-Ag(2)	59.475(16)
Ag(3)-Ag(4)	3.1767(7)	Ag(8)-Ag(1)-Ag(2)	56.253(17)
Ag(3)-Ag(8)	3.3209(8)	Ag(3)-Ag(1)-Ag(2)	61.413(15)
Ag(4)-C(21)	2.270(5)	Ag(5)-Ag(1)-Ag(2)	124.929(19)
Ag(4)-C(31)#2	2.274(5)	Ag(7)-Ag(1)-Ag(2)	2.589(6)

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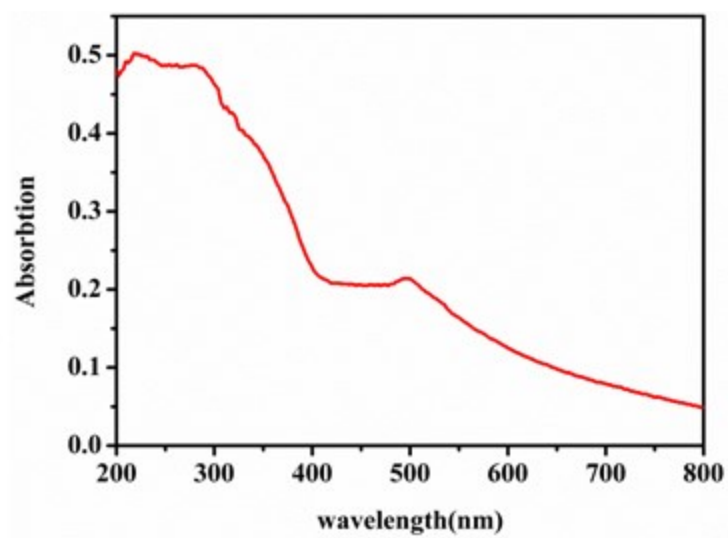


Figure S1. The solid-state UV absorption spectrum of CP₁.

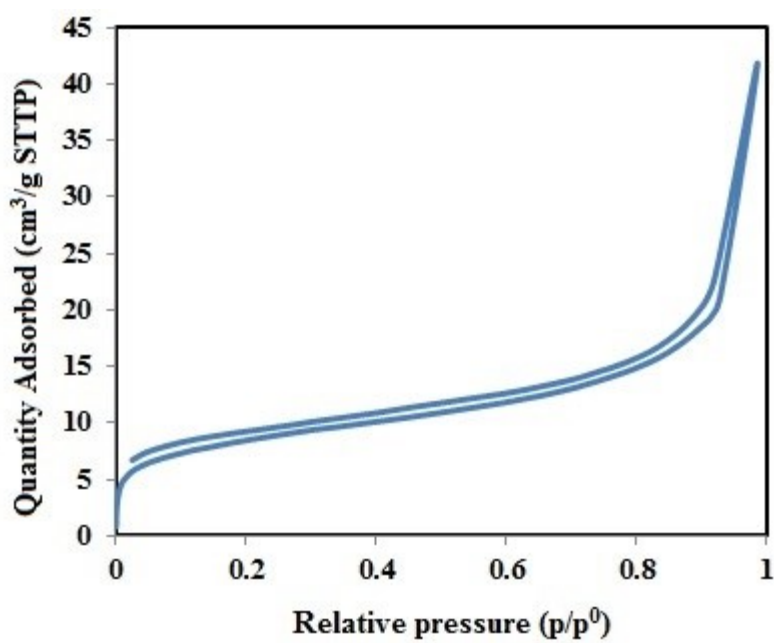


Figure S2. N₂ adsorption isotherms collected at 77 K.

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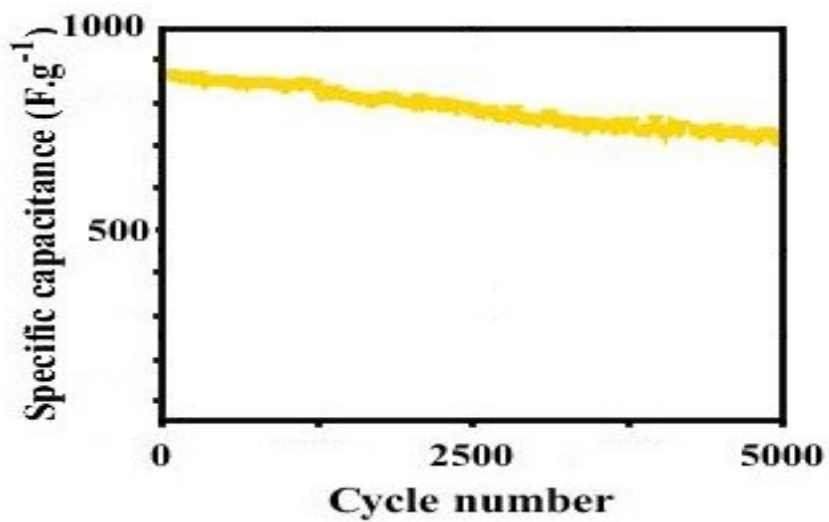


Figure S3. Cyclic stability of SSc symmetric supercapacitor at 16 A.g^{-1} for 5000 cycles.

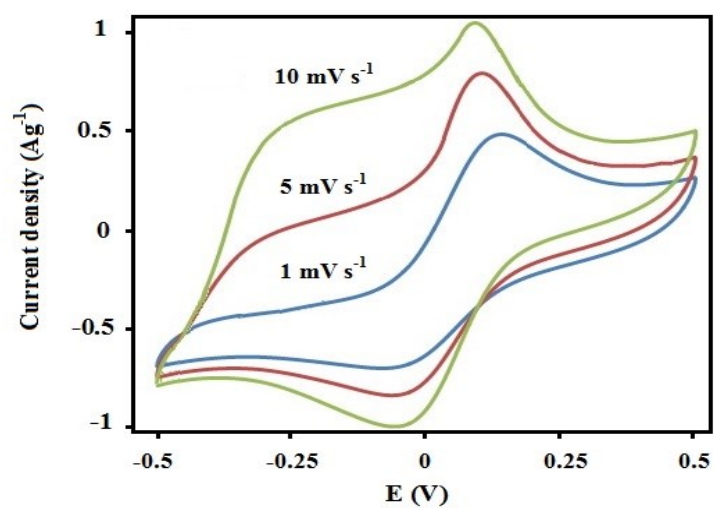


Figure S4. The CV curves of SSc at low scan rates ($1\text{-}10 \text{ mV s}^{-1}$).

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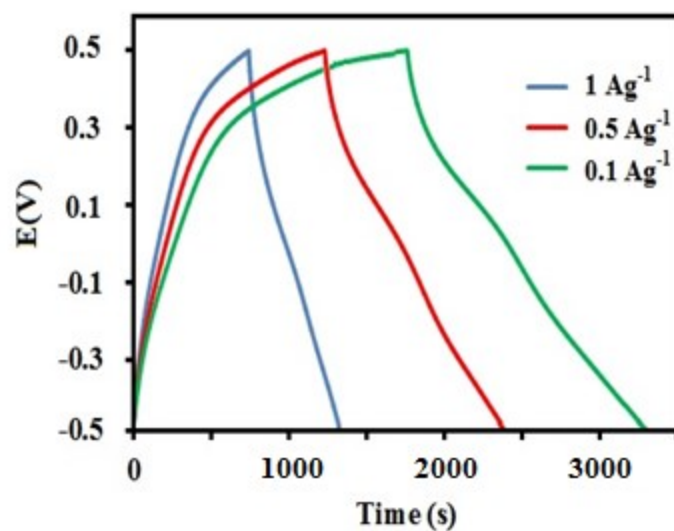


Figure S5. The charge-discharge curves at low current densities ($0.1\text{-}1 \text{ A g}^{-1}$).

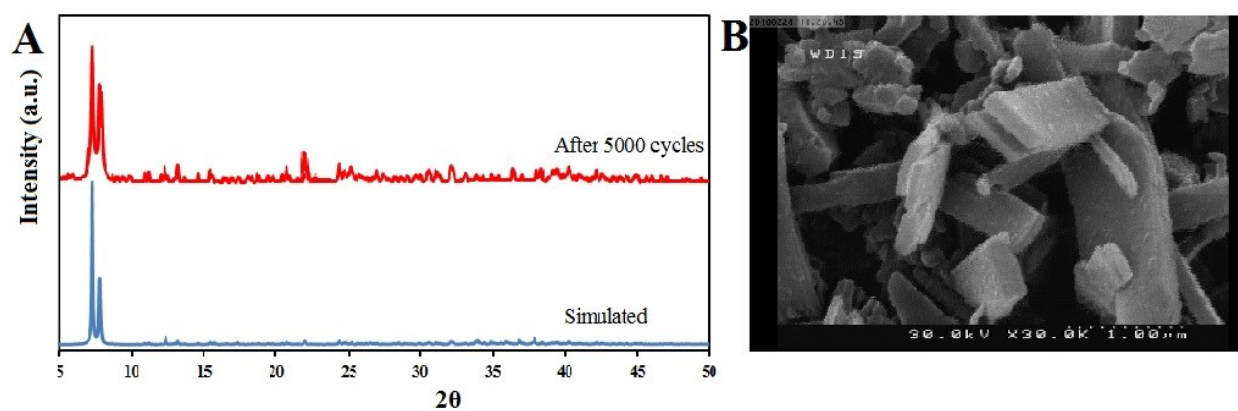


Figure S6. (A) XRD pattern of SSs after 5000 cycle in comparison with simulated one and (B) SEM images of SSs after usage as electrode.