Supporting Information

Quinolinediol molecule electrode and MXene for asymmetric supercapacitor with efficient energy storage

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Material Characterizations

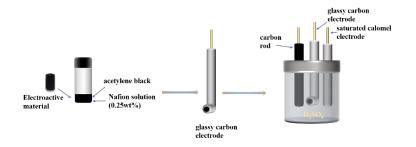
The morphology of the samples was observed by using scanning electron microscope (SEM, Ultra Plus, Carl Zeiss, Germany) and transmission electron microscope (TEM, JEM-2100 (Japan)). FTIR spectroscopy was analyzed through a Nicolet Nexus 670 instrument. The Raman spectra analysis was researched with an InVia Raman spectrometer (Renishaw) using a 514 nm argon ion laser. X-ray diffraction (XRD) analysis was generated using a diffractometer (D/Max-2400) with Cu K α radiation (λ = 1.5418 Å) at 40 kV, 100 mA. Thermogravimetric (TG) analysis was accomplished on a PerkinElmer TG/DTA-6300 instrument from 20 to 800 °C at a heating rate of 5 °C min⁻¹ in N₂ atmosphere. Surface elemental analysis was carried out with X-ray photoelectron spectroscopy (XPS) via monochromatic Al K α radiation source (ThermoVG Scientific).

Electrochemical performance test

Cyclic voltammetry (CV), galvanostatic charge/discharge (GCD) and electrochemical impedance spectroscopy (EIS) were performed on a CHI760e electrochemical workstation (Chenghua, Shanghai, China) at room temperature. LANHECT2001A testing system (Wuhan LAND electronics Co., Ltd) is used for cycling stability test.

The fabrication and electrochemical measurement of working electrodes

The fabrication of working electrodes: electrode materials (4 mg) and acetylene black (0.7mg) were uniformly dispersed in Nafion solution (0.25wt%, 0.4 mL). The above dispersion (6 μ L) was dripped onto the glassy carbon electrode by pipet gun. The electrochemical performances of electrode materials are tested in three electrode cell, which employs the glassy carbon electrode as the fluid collector, saturated calomel electrode (SCE) as the reference electrode, carbon rod as the counter electrode and 1 M H₂SO₄ solution as electrolyte (Scheme S1).



Scheme S1. The preparation process of electrode.

Calculations of capacitive characteristics

The specific capacitance (C, F g^{-1}) of active materials or asymmetric supercapacitor from the CV curves was calculated, according to the following equation:

$$C = \frac{\int IdV}{mv\Delta V} \tag{S1}$$

where I, ΔV , *v* and m are the discharging current, the potential window, the potential scan rate and the mass of active material, respectively.

To construct the asymmetric supercapacitors, the masses of the positive and negative electrodes were balanced using the following formula:

$$\frac{m_+}{m_-} = \frac{C_- \times \Delta V_-}{C_+ \times \Delta V_+} \tag{S2}$$

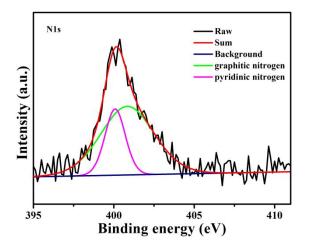
Where m, C, ΔV with positive and negative signs are the mass, capacitance of single electrodes, and applied potential range of the positive and negative electrodes, respectively.

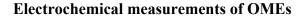
The energy density (Wh kg^{-1}) and power density (kW kg^{-1}) of the ASC device can be calculated according to equations:

$$E = \frac{I\int V(t)dt}{m}$$
(S3)
$$P = \frac{E}{t}$$
(S4)

where I is the discharging current, t is discharging time, V(t)dt is the integral area of the discharge curve and m is the mass of the active material.

Positive materials characterization





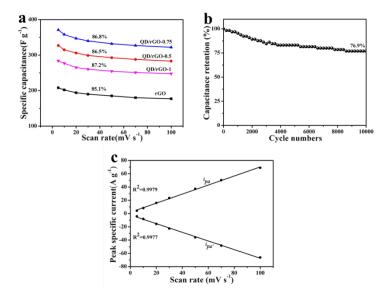


Figure S2. (a) Specific capacitance of all samples at 5–100 mV s⁻¹. (b) Cycling stability of QD/rGO-0.75. (c) Relationship of peak current and scan rate

Sample	Rct	Rs
rGO	0.34	1.99
QD/rGO-0.5	0.40	1.89
QD/rGO-0.75	0.45	1.80
QD/rGO-1	0.51	1.87

 Table S1. The Rct and Rs values of the samples

Negative materials characterizations

The microstructure of $Ti_3C_2T_x$ is studied by SEM (Figure S3a). The sample shows two dimensional sheet structure. Figure S3b shows the TEM image of $Ti_3C_2T_x$ shows thin and transparent sheets.^[1] Figure S3c shows the XRD pattern of $Ti_3C_2T_x$. A sharp peak at 6.3° and weak peaks could be observed at 13.3°, 19.1°, 26.5°, 40.6° and 60.9° corresponding to the (002), (004), (006), (008), (0012) and (110) planes, suggesting that $Ti_3C_2T_x$ is successfully synthesized.^[2] Moreover, the XPS spectrum for $Ti_3C_2T_x$ shows Ti2p, C1s, N1s, O1s and F1s peaks (Figure S3d). The high-resolution Ti2p spectrum of $Ti_3C_2T_x$ shows three doublets (Ti2p3/2-Ti2p1/2) (Figure S3e). The highresolution C1s spectrum shows five binding energies of 281.7, 282.5, 284.8, 286.2, and 288.3 eV, corresponding to C-Ti, C-Ti-O, C-C, C-O, and O-C=O (Figure S3f). The XPS results are consistent with other reports,^[3,4] further indicating that $Ti_3C_2T_x$ is successfully synthesized.

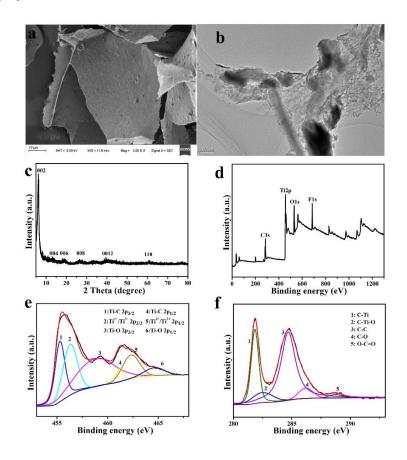


Figure S3. (a) SEM of $Ti_3C_2T_x$. (b) TEM of $Ti_3C_2T_x$. (c) XRD of $Ti_3C_2T_x$. (d) XPS spectra of $Ti_3C_2T_x$. (e) High-resolution Ti2p spectrum. (f) High-resolution C1s

spectrum.

Electrochemical measurements of $Ti_3C_2T_x$

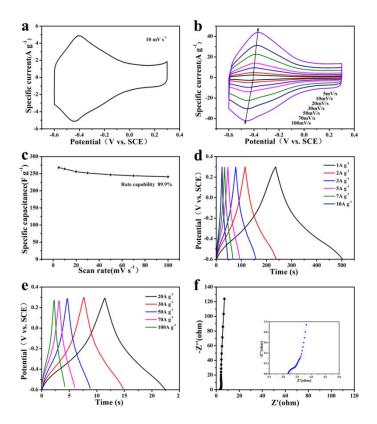


Figure S4. (a) CV curves of Ti₃C₂T_x at 10 mV s⁻¹. (b) CV curves of Ti₃C₂T_x at scan rate of 5–100 mV s⁻¹ (c) Specific capacitance of Ti₃C₂T_x at scan rate of 5–100 mV s⁻¹. (d, e) GCD curves of Ti₃C₂T_x at 1–100 A g⁻¹. (f) Nyquist plots of Ti₃C₂T_x (The inset in (f) shows the magnified high-frequency region).

Electrochemical measurement of ASC

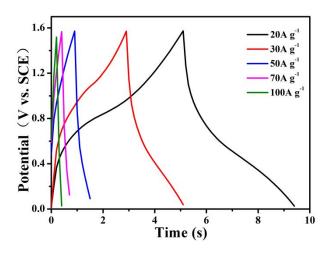


Figure S5. GCD curves of ASC at $20-100 \text{ A g}^{-1}$.

References

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