Support Information

NiCoP/MXene nanocomposites via electrostatic self-assembly for

high-performance supercapacitor electrodes

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1. Electrochemical Measurements

The specific capacity (Cs, C g⁻¹) of the electrode material can be obtained from the galvanostatic charge-discharge curve by the following formula:¹

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

Where *I* is the discharge current (A), Δt is the discharge time (s), and *m* represents the mass of the active material coated on the carbon cloth (g).

Asymmetric supercapacitor (ASC) was assembled using NiCoP/MXene composites and commercial activated carbon (AC) as positive electrode and negative electrode. In order to balance the potential of the two electrodes, the mass ratio of the positive and negative materials must satisfy the following formula:²

$$\frac{m_+}{m_-} = \frac{C_-\Delta V_-}{C_+\Delta V_+} \tag{2}$$

Where m (g) is the mass of the electrode active material, C (C g⁻¹) is the specific capacitance of a single electrode, ΔV is the potential window of the positive electrode and the negative electrode, and the mass ratio of the positive electrode and the negative electrode is calculated to be 1:5.24. The corresponding energy density (E, Wh kg⁻¹) and power density (P, W kg⁻¹) can be obtained by the following two formulas:³

$$E = \frac{C \times \Delta V^2}{2 \times 3.6} \tag{3}$$

$$P = \frac{3600 \times E}{\Delta t} \tag{4}$$

2. Material characterization

The phase structure of the NiCoP/MXene composite was characterized by Bruker D8 advanced X-ray powder diffractometer with Cu Ka radiation ($\lambda = 1.5418$ Å). The surface morphology and energy spectrum (EDS) of the samples were analyzed by field emission scanning electron microscopy (SEM, FE-SEM, Hitachi S-4800, Japan). The microstructure of the samples was characterized using transmission electron

microscopy (TEM, FEI Tecnai G²F20, USA). The elemental composition and valence states of the samples were analyzed by X-ray photoelectron spectroscopy (XPS, UK AXIS SUPRA (Kratos)).

3. Supplementary figures

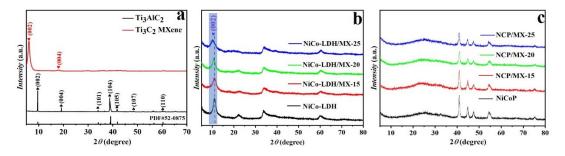


Figure S1. (a) XRD patterns of Ti_3AlC_2 and Ti_3C_2 MXene; (b, c) XRD patterns of NiCo-LDH precursors and NiCoP with different masses of MXene nanosheets added.

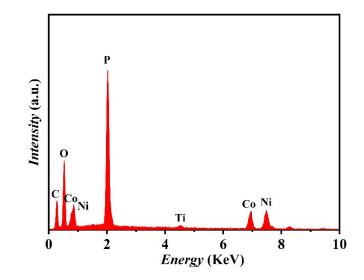


Figure S2. EDS spectrum of NCP/MX-20.

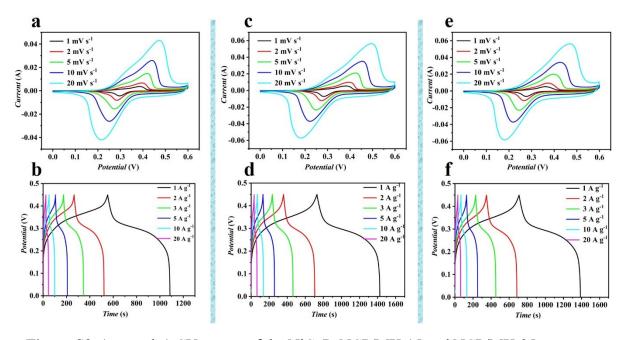


Figure S3. (a, c and e) CV curves of the NiCoP, NCP/MX-15 and NCP/MX-25 at different scan rates; (b, d and f) GCD curves of the NiCoP, NCP/MX-15 and NCP/MX-25 at different current densities.

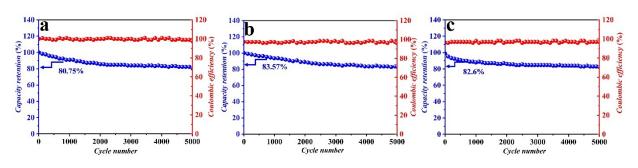


Figure S4. (a, b and c) Cycling performance of the NiCoP, NCP/MX-15 and NCP/MX-25 electrode within 5000 cycles at 10 A g⁻¹.

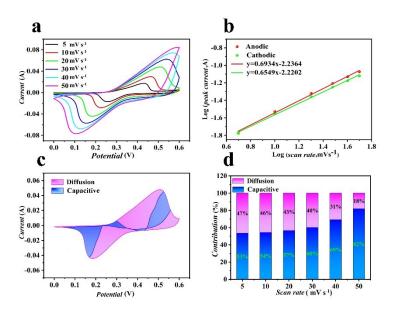


Figure S5. (a) CV curves of NiCoP electrode at different scan rates; (b) the relationship between log(v) and log(i) of NiCoP electrode at different scan rates; (c) distribution of diffusion and capacitive contribution of NiCoP electrode at 20 mV s⁻¹ and (d) contribution rate of diffusion and capacitive control behavior of NiCoP electrodes at different scan rates.

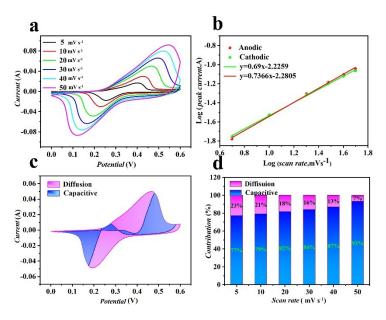


Figure S6. (a) CV curves of NCP/MX-15 electrode at different scan rates; (b) the relationship between log(v) and log(i) of NCP/MX-15 electrode at different scan rates;
(c) distribution of diffusion and capacitive contribution of NCP/MX-15 electrode at 20 mV s⁻¹ and (d) contribution rate of diffusion and capacitive control behavior of NCP/MX-15 electrodes at different scan rates.

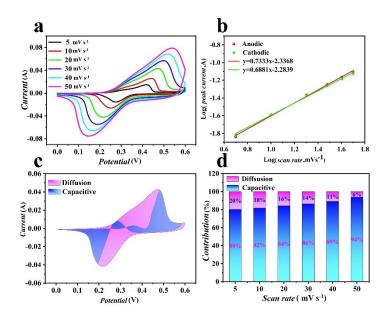


Figure S7. (a) CV curves of NCP/MX-25 electrode at different scan rates; (b) the relationship between log(v) and log(i) of NCP/MX-25 electrode at different scan rates;
(c) distribution of diffusion and capacitive contribution of NCP/MX-25 electrode at 20 mV s⁻¹ and (d) contribution rate of diffusion and capacitive control behavior of NCP/MX-25 electrodes at different scan rates.

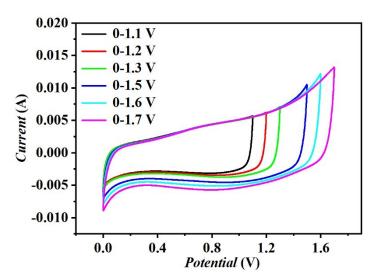


Figure S8. CV curves of the NCP/MX-20//AC ASC at different voltage windows at a scan rate of 20 mV s⁻¹.

4. Supplementary tables S1-S4

Table S1. The NCP/MX-20 of elemental composition of Smart quant results by theEDS.

Element	Wt%	At%
РК	45.57	61.23
NiK	30.54	21.66
СоК	22.49	15.89
TiK	1.4	1.22

Table S2. The specific capacititance (C g^{-1}) at different current densities in Figure 3e.

	1 A g ⁻¹	2 A g ⁻¹	3 A g ⁻¹	5 A g ⁻¹	10 A g ⁻¹	20 A g ⁻¹
NiCoP	532.67	510.42	509	508.53	482.99	444
NCP/MX-15	696.83	691.32	683.74	672.55	652	618
NCP/MX-20	848.83	844.69	842.69	830.05	807.02	764
NCP/MX-25	677.93	659.14	656.94	645.54	622.99	582

Table S3. Comparison of the capacity performance of as-prepared NCP/MX-20 and

 battery-type electrode materials reported previously.

	1			
Electrode material	Morphology	Specific	Current	Ref
		capacity	density	
NiCoP/C	nanoparticles	775.87 C g^{-1}	1 A g ⁻¹	4
NiCoP/NC	hollow	1127 F g^{-1}	1 A g ⁻¹	5
	nanocages			
NiCoP@CoS	tree-like core-	$1796 \ F \ g^{-1}$	2 A g^{-1}	6
	shell			
	nanoarrays			
Cu-NiCoP	nanocages	$800 \ C \ g^{-1}$	$1 \mathrm{A} \mathrm{g}^{-1}$	7
NiCoP/NiCo-OH	hierarchical	1100 F g ⁻¹	1 A g ⁻¹	8
	cactus-like	5	C	

Ni-Co-P/POx	Sea-urchin-like	$647 \ C \ g^{-1}$	$1 \mathrm{A} \mathrm{g}^{-1}$	9
NiCoP/MLG	porous flower-	1419.6 F g ⁻¹	$1 \mathrm{A} \mathrm{g}^{-1}$	10
	like			
N-	Hetero-	152 mAh g ⁻¹	$1 \mathrm{A} \mathrm{g}^{-1}$	11
CNTs@NiCoP/CoP	nanosheets	(547.2 C g ⁻¹)		
NiCoP	porous	1142.84 F g ⁻¹	1 A g ⁻¹	12
	nanosheet			
LDH-MXene	nanosheets	983.6 F g ⁻¹	2 A g ⁻¹	13
NiCoP	nanoparticles	1456 F g ⁻¹	2 A g ⁻¹	14
Ni-CoP@C@CNT	nanorods	708.1 F g ⁻¹	1 A g ⁻¹	15
NiCoP-4-500	hollow nano	1590 F g ⁻¹	1 A g ⁻¹	16
	cubes			
O-NiCoP	porous	1663.2 F g^{-1}	1 A g ⁻¹	17
	nanosheets			
NiCoP-MOF	lamellar brick-	728 C g ⁻¹	1 A g ⁻¹	18
	stacked			
Ni ₂ P / NiCoP	nanoparticles	741.3 C g ⁻¹	1 A g ⁻¹	19
NCP/MX-20	porous flower-	848.8 C g ⁻¹	1 A g ⁻¹	This work
	like	(1886.2 Fg ⁻¹)		

Table S4. Fitted parameters of the Nyquist plots in Fig. 3f using equivalent circuits.

samples	$R_{s}\left(\Omega ight)$	$R_{ct}\left(\Omega ight)$
NiCoP	1.125	0.23
NCP/MX-15	0.664	0.12
NCP/MX-20	0.46	0.13
NCP/MX-25	0.71	0.19

5. Supplementary References

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