

Supporting information

Engineering highly active surfaces in molybdenum-integrated cobalt telluride electrodes for enhanced battery-type charge-storage in supercapacitors

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S1. Preparation of CoTe₂ nanostructures

Sulfuric acid-treated TeO₂ (1.5 mM) was mixed with an aqueous (NH₄)₂Co(SO₄)₂ solution (2 mM) under vigorous magnetic stirring at room temperature for 1 h. Subsequently, 2 mL of N₂H₄ and 4 mL of NH₄OH were slowly introduced under continuous stirring to obtain a homogeneous reaction mixture. The mixture was then heated on a hot plate at 80 °C for 3 h while maintaining constant agitation. After cooling naturally to room temperature, the resulting precipitate was collected and washed several times with DI water and ethanol via centrifugation. The black CoTe₂ powder was then dehydrated at 60 °C for 3 h in a hot-air oven and finally calcined in a tellurium atmosphere at 500 °C for 2 h.

S2. Preparation of MoTe₂ nanostructures

Sulfuric acid-treated TeO₂ (1.5 mM) was mixed with a (NH₄)₆Mo₇O₂₄ solution (2 mM) under vigorous magnetic stirring at room temperature for 1 h. Afterwards, 2 mL of N₂H₄ and 2 mL of NH₄OH were slowly added under continuous magnetic stirring to form a uniform solution mixture. The reaction mixture was then heated at 80 °C for 3 h with constant stirring. After gradually cooling to room temperature, the resulting precipitate was collected and rinsed with DI water and ethanol by centrifugation. The black MoTe₂ powder was dehydrated at 60 °C for 3 h and then calcined in a tellurium environment at 500 °C for 2 h.

S3. Characterisations

The physical properties of the synthesized Co_xTe₂-based nanostructures were analysed using the following instruments: JEOL JSM-6700F field-emission scanning electron microscope; JEOL-2010F transmission electron microscope; Microtrac BELsorp-mini II system at 77 K for BET analysis; Renishaw inVia RE04 Raman spectrometer; and PHI 5000 VersaProbe II X-ray photoelectron spectrometer.

Mass calculation:

The mass loading of AC in the asymmetric device was calculated based on equation S1.

$$\frac{m_+}{m_-} = \frac{C_{sp-} \times \Delta V_-}{C_{sp+} \times \Delta V_+} \quad (\text{S1})$$

where m_- and m_+ represent the mass of active materials, C_s^- and C_s^+ represent specific capacity, ΔV_- and ΔV_+ represent the potential windows for the negative and positive electrodes, respectively.

The ASC device-specific capacitance (C_{SC}) was estimated using equation (S2)

$$C_{SC} = \frac{i(\Delta t)}{w(\Delta V)} \quad (\text{S2})$$

The energy density and power density parameters of ASCs were obtained via the following relationships.

$$E = \frac{C_{SC} \times (\Delta V)^2}{2 \times 3.6} \quad (\text{S3})$$

$$P = \frac{E}{\Delta t} \quad (\text{S4})$$

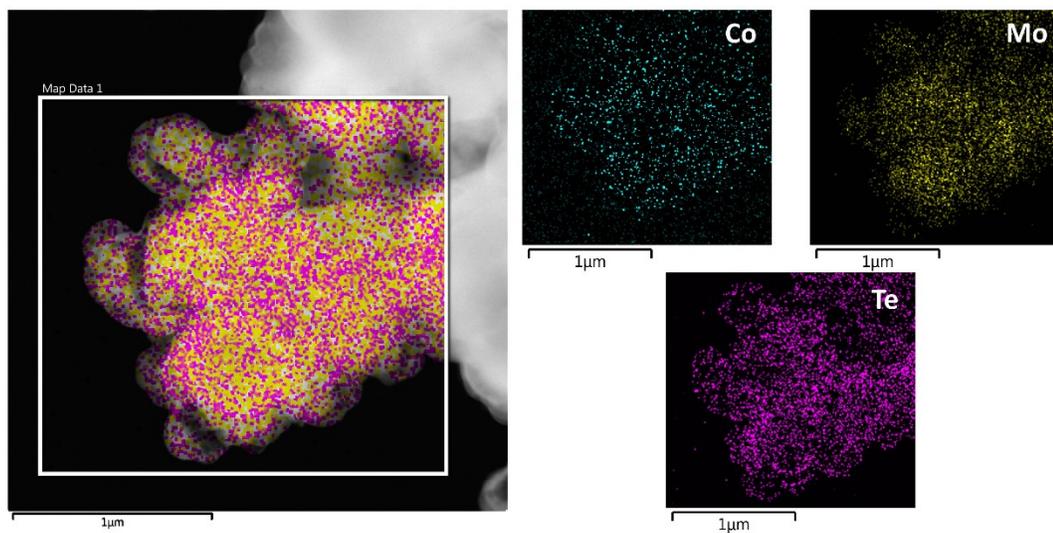


Figure S1. TEM elemental mapping of MCT-1 and corresponding distributions of Co, Mo, and Te.

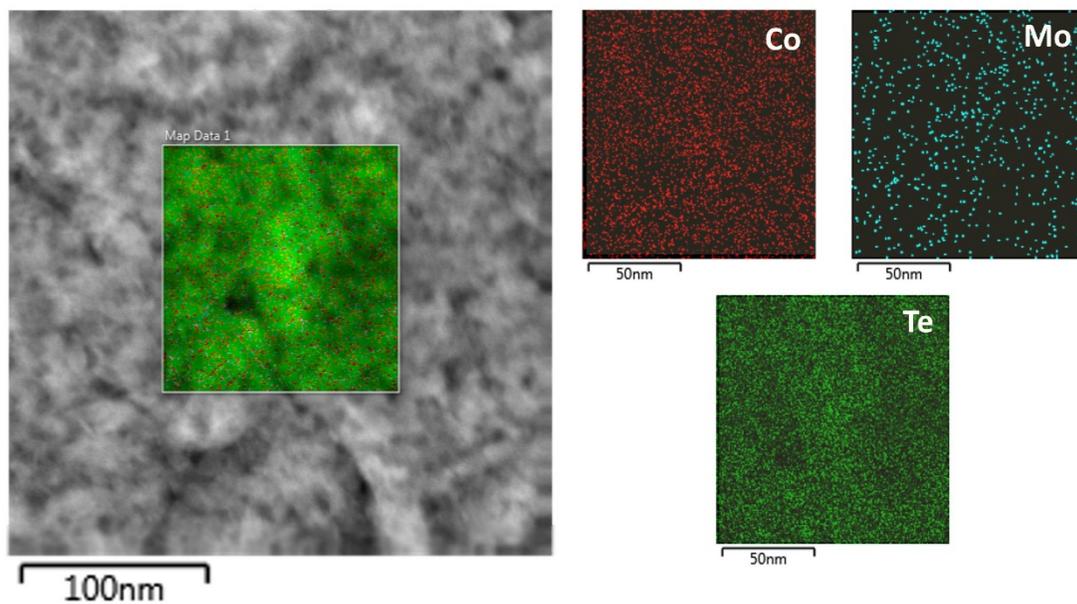


Figure S2. TEM elemental mapping of MCT-2 and corresponding distributions of Co, Mo, and Te.

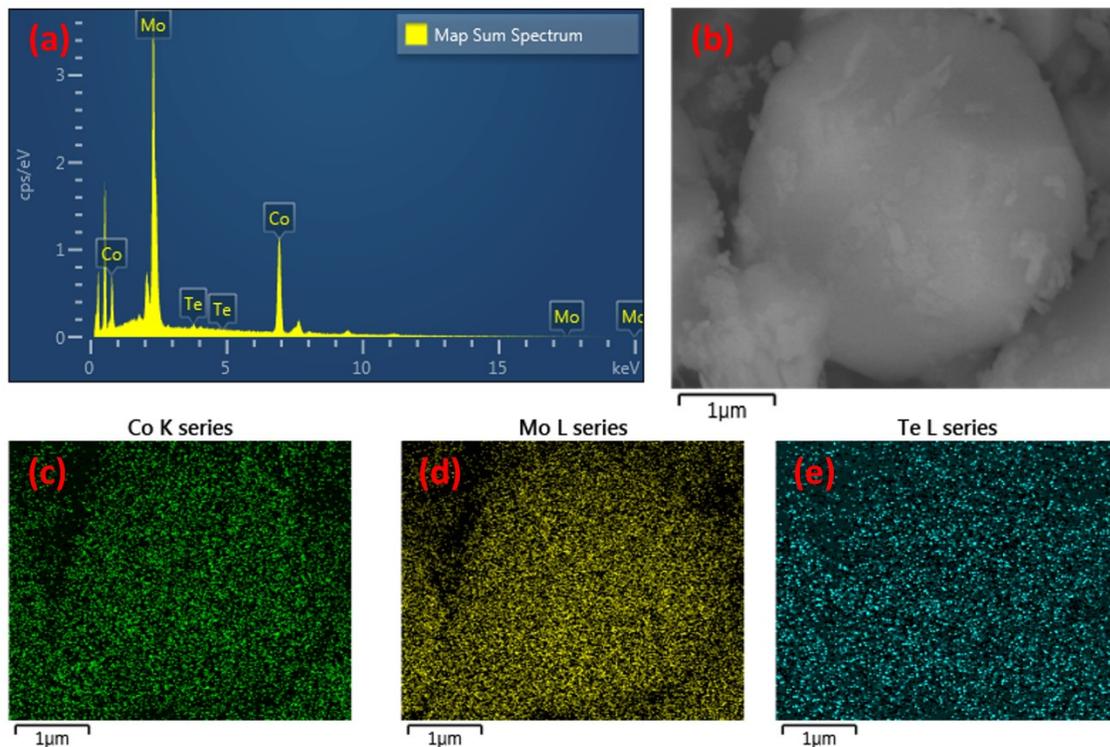


Figure S3. (a) EDS spectrum of MCT-3; elemental mapping of MCT-3 shown by (b) SEM mapping and corresponding distributions of (c) Co, (d) Mo, and (e) Te.

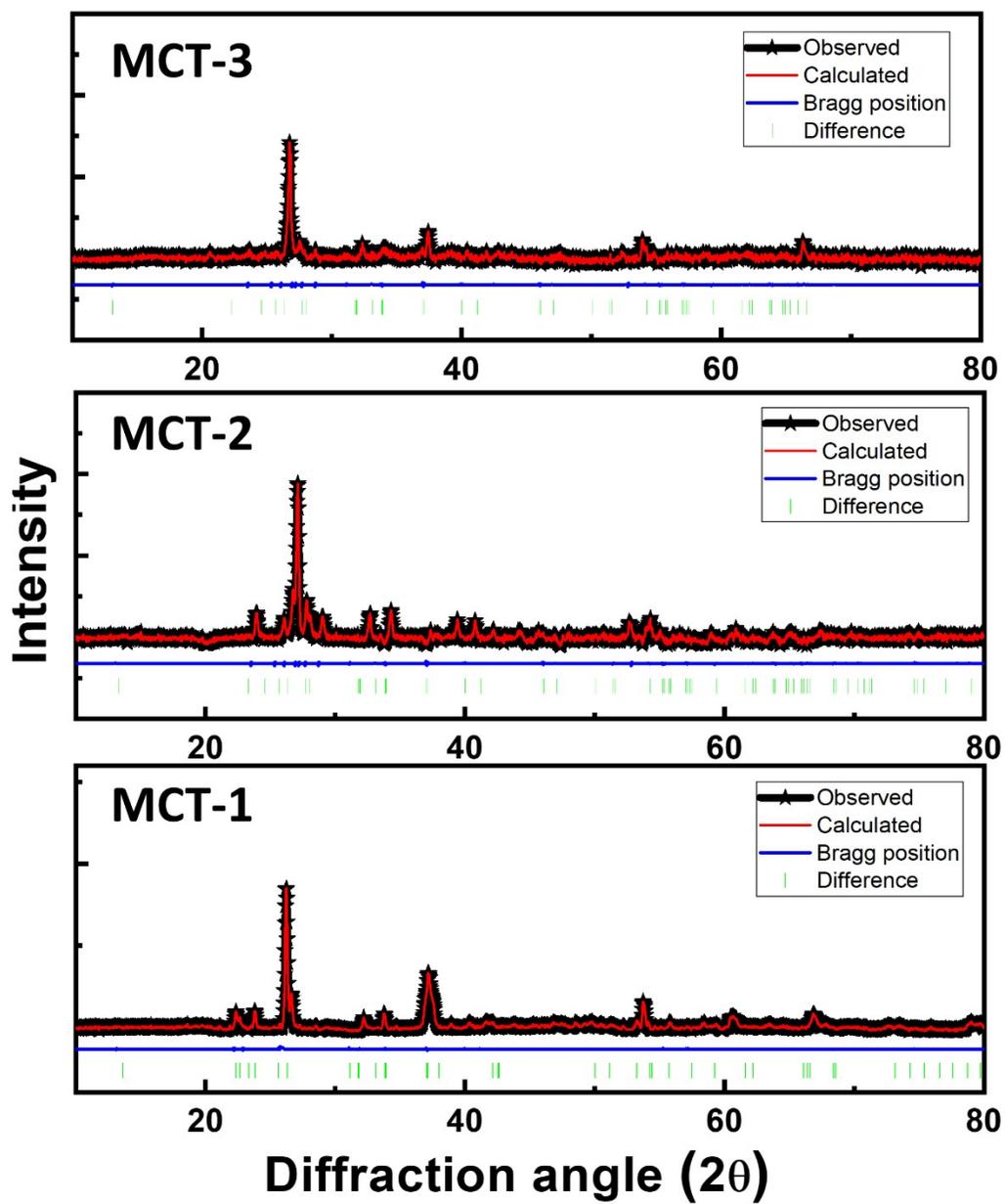


Figure S4. Rietveld refinement on MCT-1, MCT-2, and MCT-3 patterns.

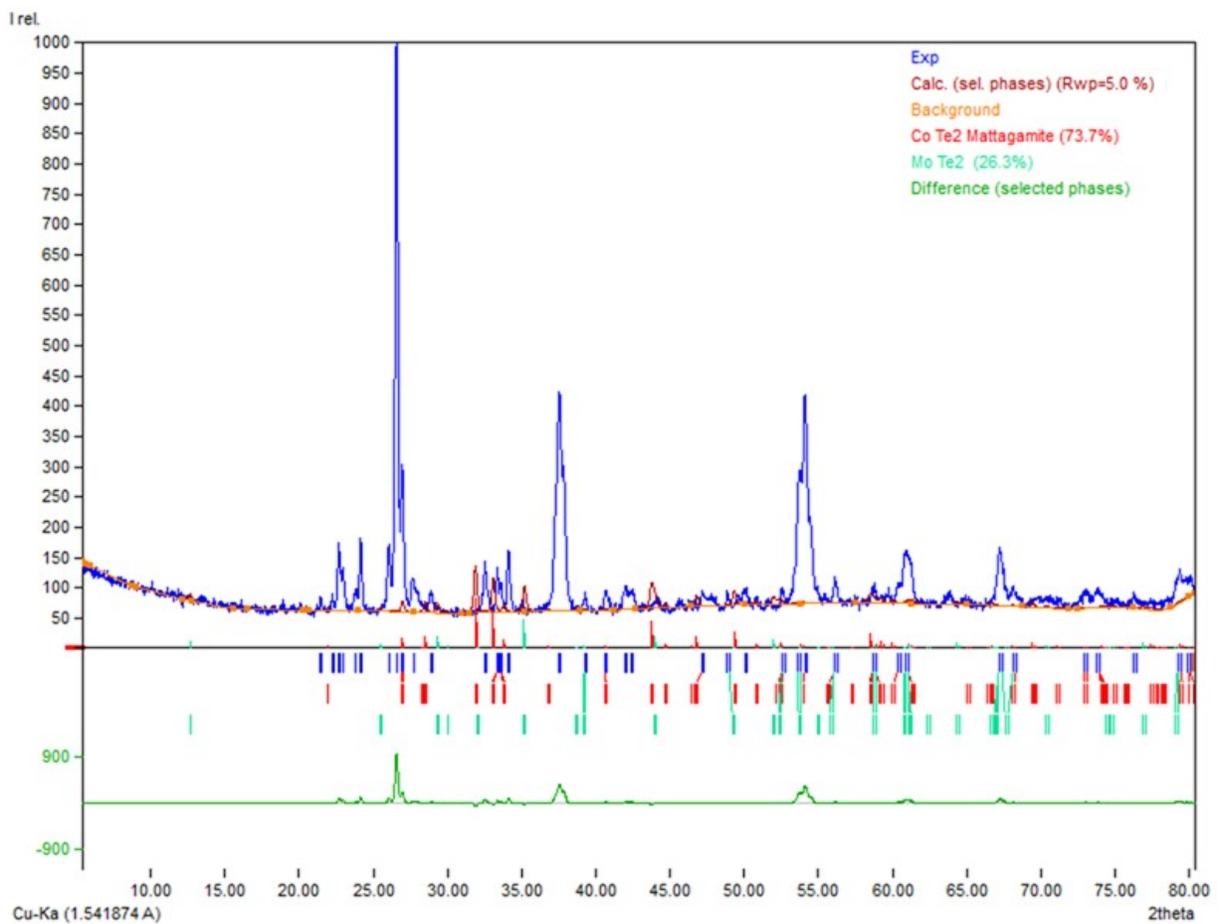


Figure S5. Phase composition ratio of MCT-1 solid solution.

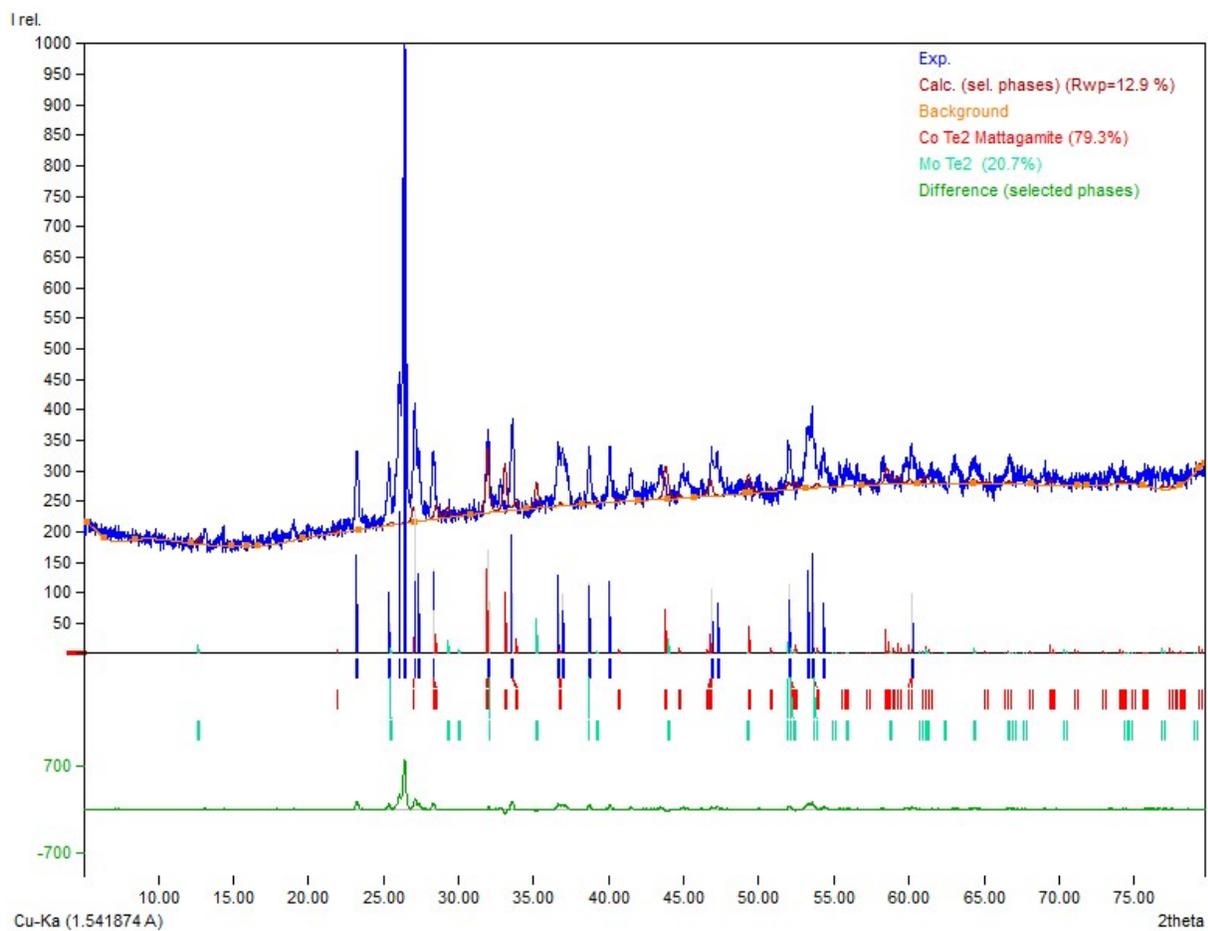


Figure S6. Phase composition ratio of MCT-2 solid solution.

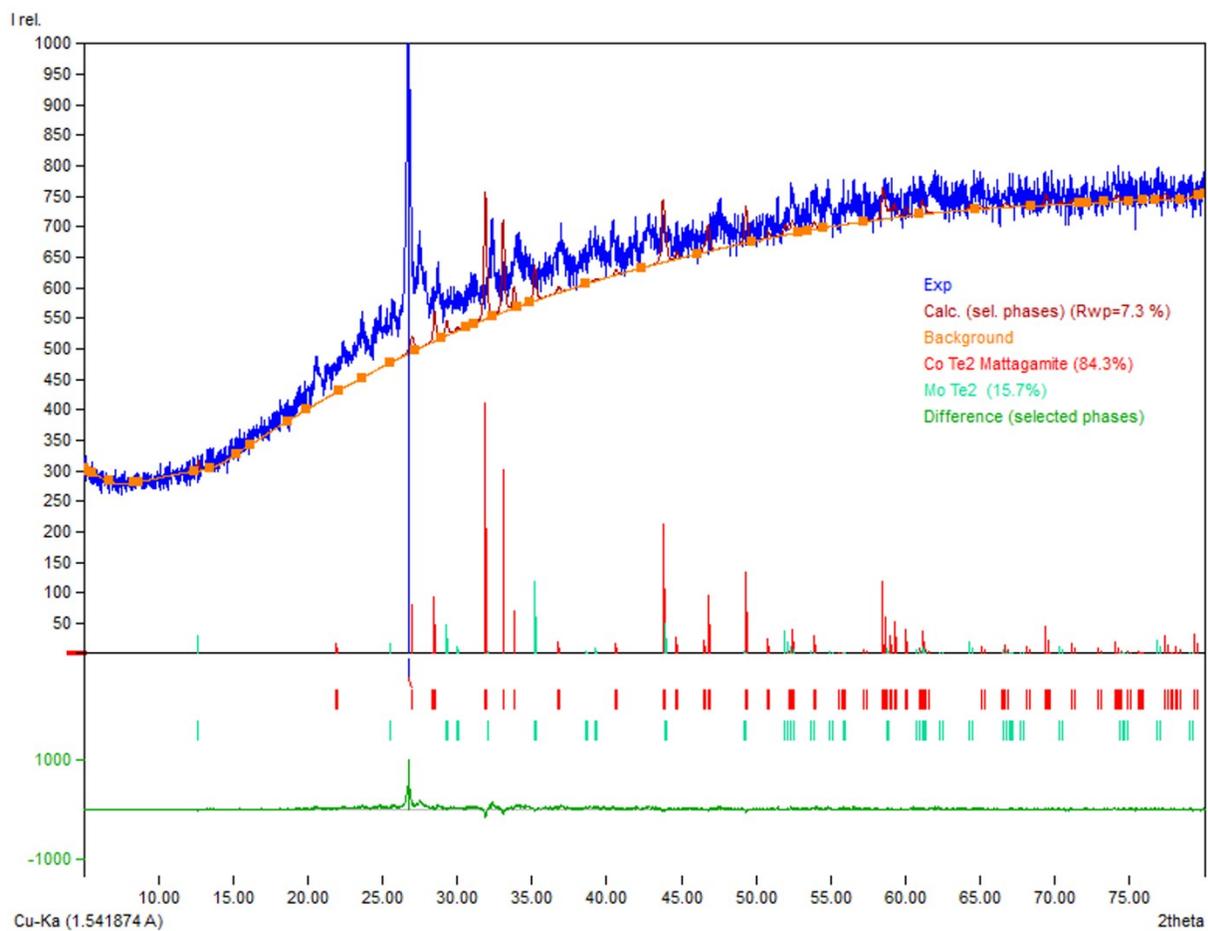


Figure S7. Phase composition ratio of MCT-3 solid solution.

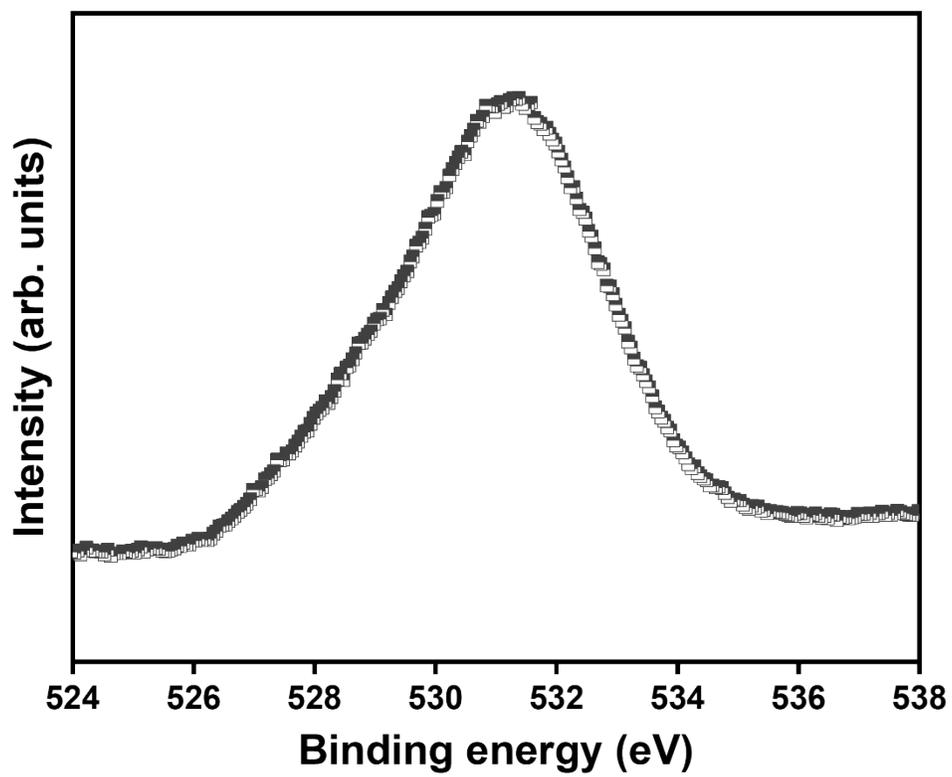


Figure S8. O 1s XPS profile for MCT-3 structure.

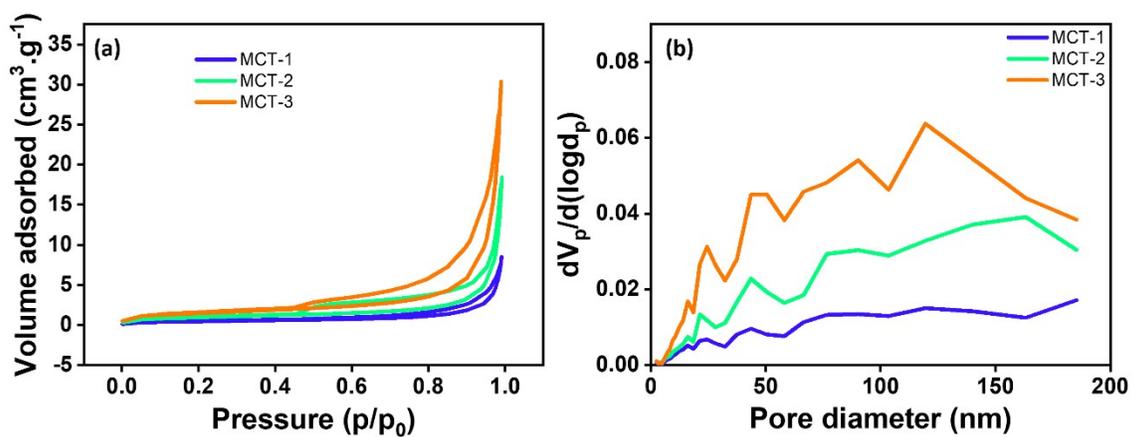


Figure S9. (a) BET surface-area profiles and (b) BJH pore-size distribution curves for MCT-1, MCT-2, and MCT-3 nanostructures.

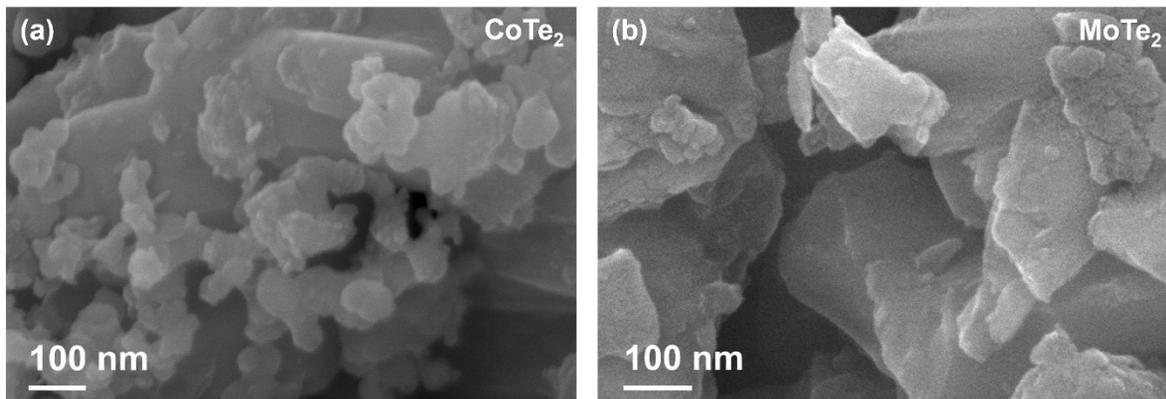


Figure S10. SEM of (a) CoTe₂ and (b) MoTe₂ nanostructures.

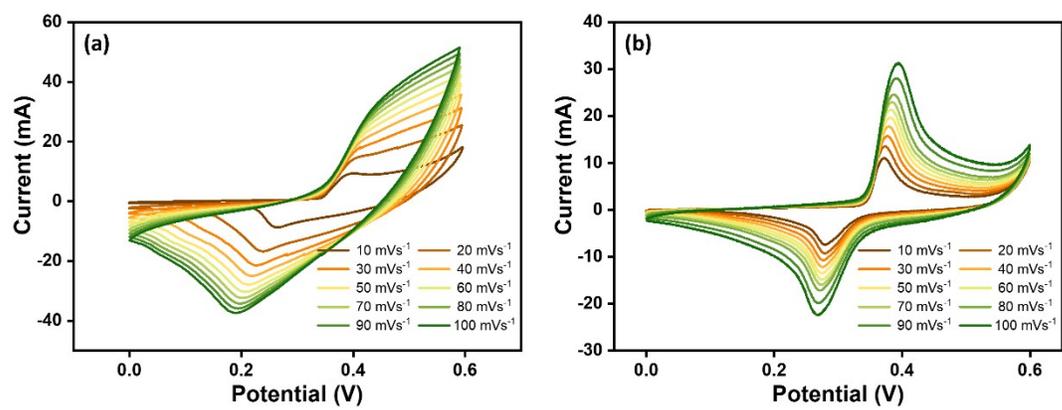


Figure S11. CV profiles of (a) CoTe₂ and (b) MoTe₂ nanostructures recorded at various scan rates.

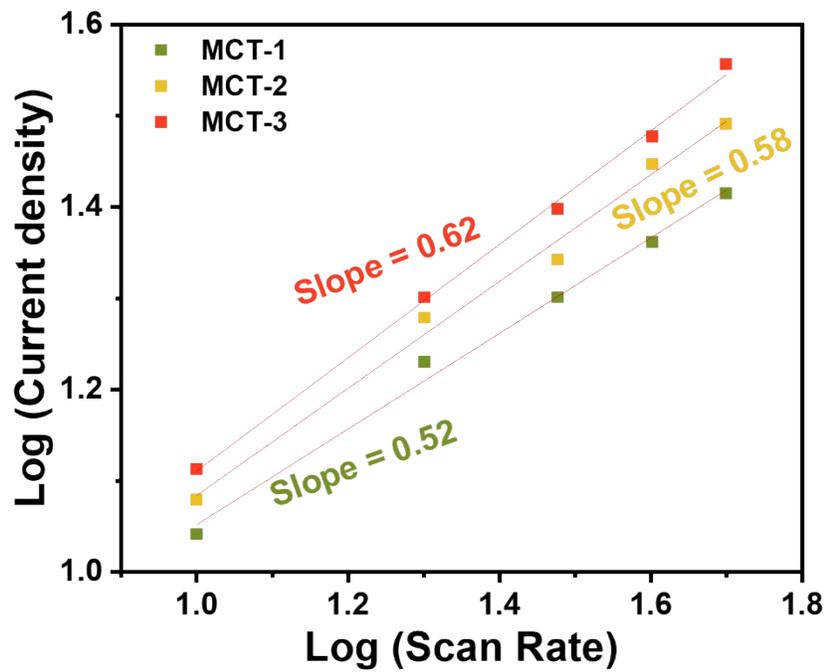


Figure S12. Plot of current density vs scan rate for MCT-1, MCT-2, and MCT-3 nanostructures.

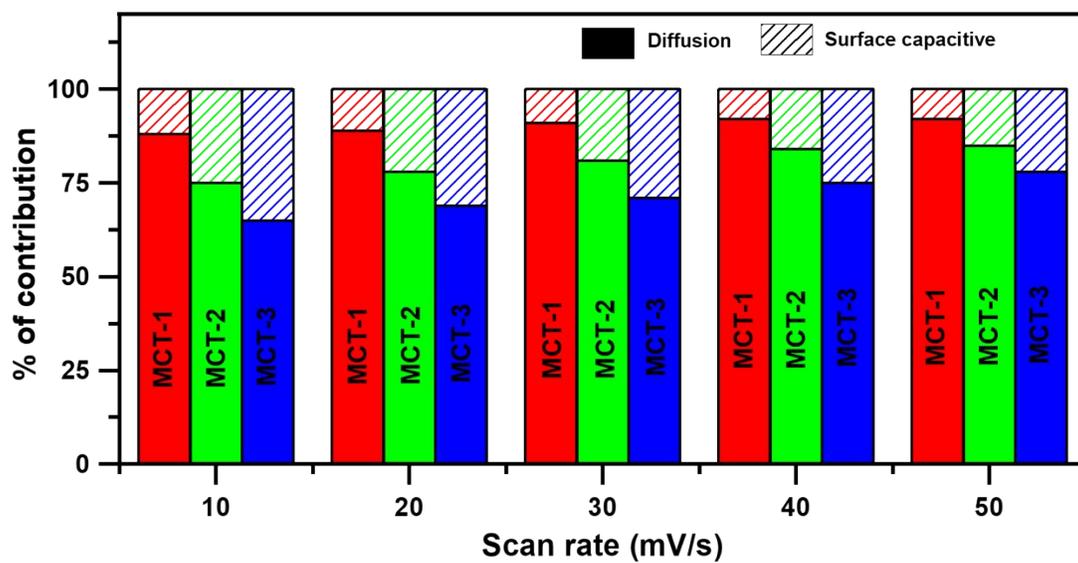


Figure S13. The capacitive and diffusive charge contribution in the MCT-1, MCT-2, and MCT-3 nanostructures electrodes as a function of the scan rate.

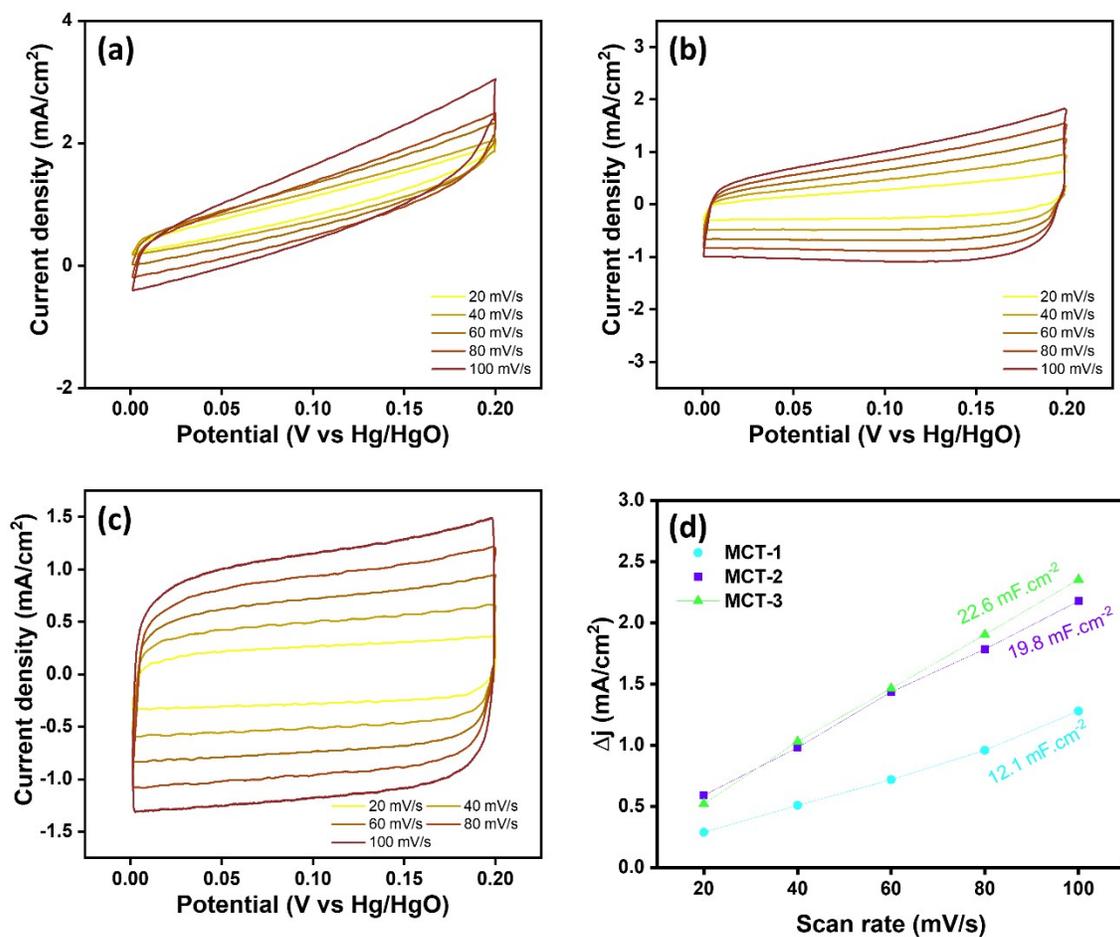


Figure S14. Non-faradaic region CVs at different scan rates for the (a) MCT-1, (b) MCT-2, and (c) MCT-3 nanostructures; current density variations at 0.1 V vs Hg/HgO for MCT-1, MCT-2, and MCT-3 nanostructures.

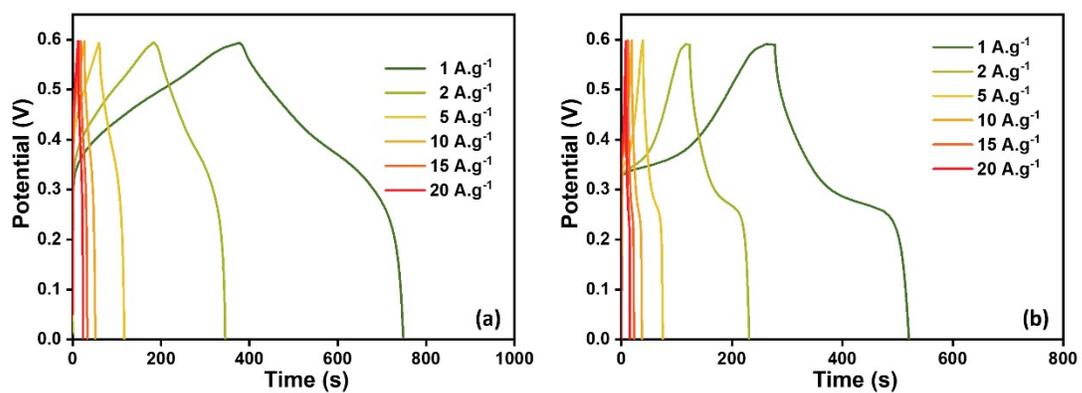


Figure S15. GCD profiles of (a) CoTe₂ and (b) MoTe₂ nanostructures at different applied current densities.

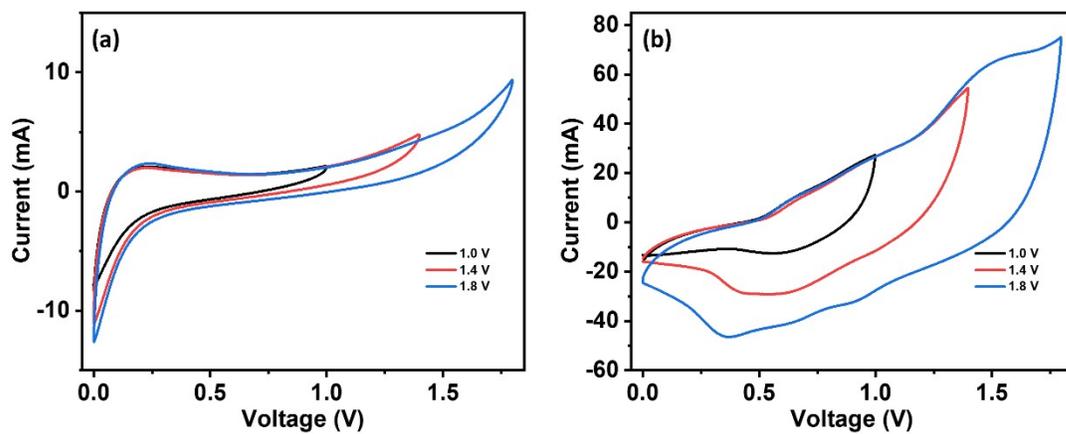


Figure S16. CV profiles of (a) MCT-1//AC and (b) MCT-2//AC asymmetric devices at different voltage windows.

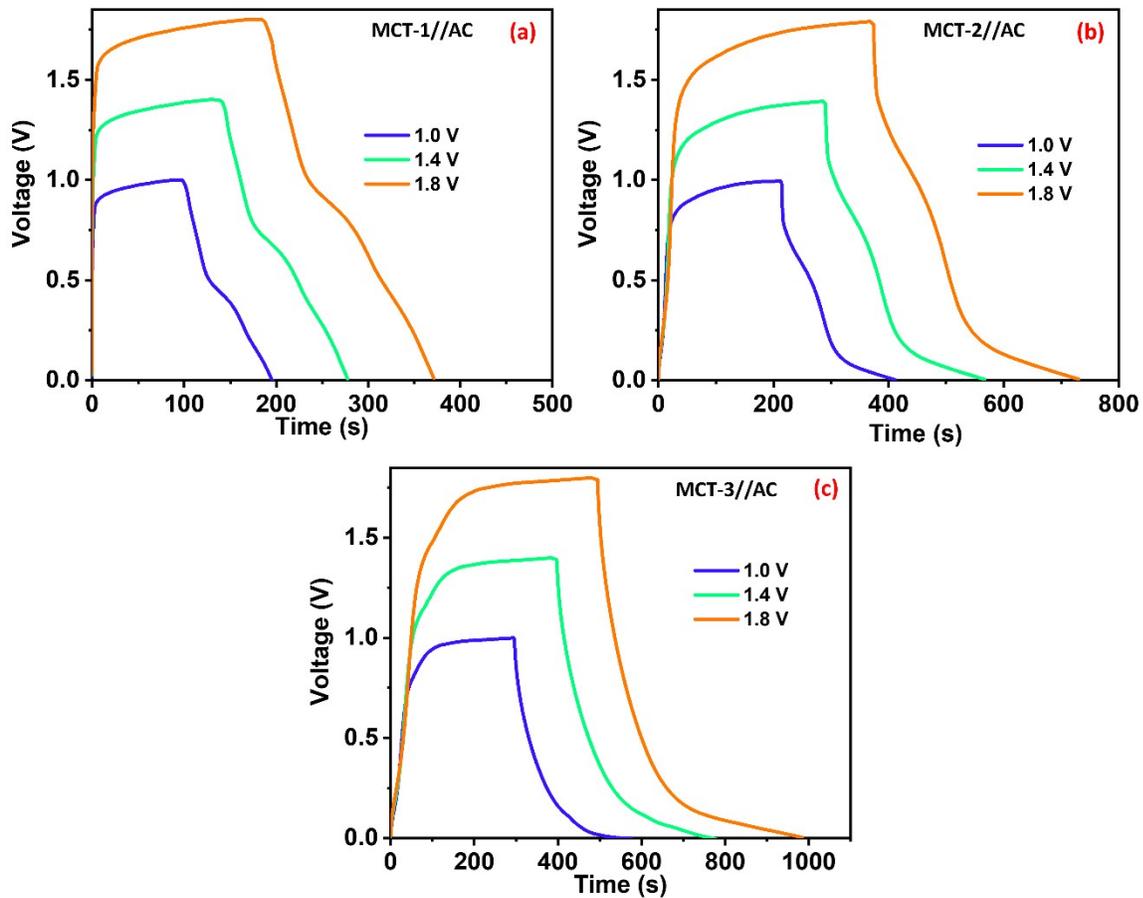


Figure S17. GCD profiles of (a) MCT-1//AC, (b) MCT-2//AC and (c) MCT-3//AC asymmetric devices at different voltage windows.

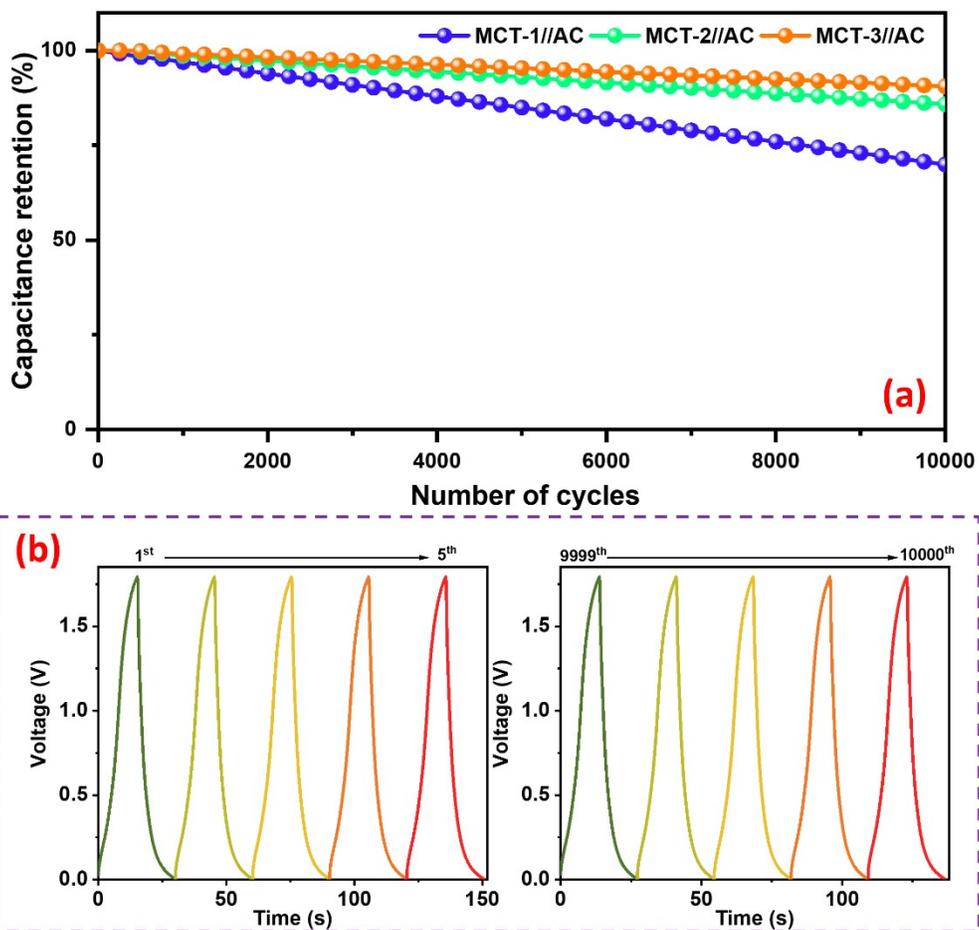


Figure S18. (a) Capacity retention of MCT-1, MCT-2 and MCT-3 at 20 A g^{-1} ; (b) initial and final five GCD cycles for MCT-3.

Table S1. N₂ sorption analysis parameters for MCT-1, MCT-2, and MCT-3 nanostructures.

Sample	BET (m².g⁻¹)	Pore volume (cm³.g⁻¹)	Mean pore diameter (nm)
MCT-1	1.68	0.012	29.2
MCT-2	3.51	0.026	29.7
MCT-3	5.21	0.046	35.7

Table S2. Three-electrode supercapacitor performance comparison with TMDs electrodes.

Electrode materials	Electrolyte (M)	Specific capacitance	Capacitance retention (%) / cycles	Ref.
<i>MCT-3</i>	<i>1 KOH</i>	<i>691 C·g⁻¹ (1151 F·g⁻¹) @ 1 A·g⁻¹</i>	<i>98 / 10000</i>	<i>In this study</i>
CoTe	3 M KOH	622.8 F·g ⁻¹ @ 1 A·g ⁻¹	85 / 2000	1
NiTe ₂	3 M KOH	804 F g ⁻¹ @ 1 A g ⁻¹	91.3 / 1000	2
NiTe/ZIF-67	3 M KOH	1521 F·g ⁻¹ @ 1A·g ⁻¹	93.5/5000	3
MoTe ₂ /graphene	1 M Na ₂ SO ₄	434 ± 37.5 F g ⁻¹ @ 1A g ⁻¹	-	4
MoTe ₂	1.0 M KOH	416 F·g ⁻¹ @ 2 A·g ⁻¹	97.1% / 5000	5
CuTe-CoTe@ NiTe ₂	6 M KOH	1481 F·g ⁻¹ @ 1 A·g ⁻¹	92.15 / 10000	6
MWCNTs/MoTe ₂	1 M NaOH	502 F g ⁻¹ @ 2 mV s ⁻¹	81 / 5000	7
CoTe/NiCo ₂ S ₄	3 M KOH	1470.45 F g ⁻¹ @ 1 A g ⁻¹	86.69 / 5000	8
NiCoTe ₂	6 K KOH	2006 F·g ⁻¹ @ 1 A·g ⁻¹	96.15 / 20000	9
CoTe ₂ nanoflowers	1 M KOH	460 F g ⁻¹ @ 1.5 A g ⁻¹	91 / 5000	10
NiTe ₂ -Co ₂ Te ₂ @rGO	1 M KOH	223.6 mAh g ⁻¹ at 1 A g ⁻¹	89.3 / 3000	11
NG-CoTe ₂ /CRF	2 M KOH	301 F g ⁻¹ @ 1 Ag ⁻¹	98.6 / 5000	12
ZnFe ₂ O ₄ -rGO	2 M KOH	352.9 F g ⁻¹ @1 A g ⁻¹	92/10000	13
1T'-MoTe ₂ nanosheets	2 M KOH	1393 F g ⁻¹ @1 A g ⁻¹	87 (50 A g ⁻¹) / 1000	14
CoTe NRs	3.5 M KOH	170 C·g ⁻¹ @ 0.5 A·g ⁻¹	99 / 5000	15

Table S3. Two-electrode supercapacitor devices using the various TMDs electrodes.

Electrode materials	Specific capacitance	Energy density	Power density	Capacitance retention (%) / cycles	Ref.
<i>MCT-3//AC</i>	<i>271 F g⁻¹ @ 1 A g⁻¹</i>	<i>122 Wh kg⁻¹</i>	<i>2000 W kg⁻¹</i>	<i>90 / 10000</i>	<i>This work</i>
CoTe//AC	192.1 F·g ⁻¹ @ 1 A·g ⁻¹	67 Wh Kg ⁻¹	793 W kg ⁻¹	94.8 / 5000	1
NiTe//AC	94.6 F g ⁻¹	33.6 Wh. kg ⁻¹	807.1 W. kg ⁻¹	81/3000	2
NiTe/ZIF-67//AC	62.6 F·g ⁻¹ @ 0.2 A·g ⁻¹	45.3 Wh Kg ⁻¹	5689.1 W kg ⁻¹	94.8 / 5000	3
MoTe ₂ /graphene - symmetric	138 F·g ⁻¹ @ 1 A·g ⁻¹	43.2 Wh kg ⁻¹	3000 W kg ⁻¹	98/10000	4
MoTe ₂ //AC	138 F g ⁻¹ @ 2 A g ⁻¹	49 Wh kg ⁻¹	961 W kg ⁻¹	95.5 / 5000	5
Nf@CCT@NT//AC	274 F·g ⁻¹ @ 1 A·g ⁻¹	41.92 Wh kg ⁻¹	33573.7 W kg ⁻¹	89.35 / 10000	6
SS/MWCNTs/ MoTe ₂ - symmetric	68.01 @ 0.2 mA.cm ⁻²	18.51 Wh kg ⁻¹	416 W kg ⁻¹	94 / 2000	7
CoTe/NiCo ₂ S ₄ //AC	88.89 F g ⁻¹ @ 1 A g ⁻¹	39.8 Wh kg ⁻¹	832 W kg ⁻¹	86.61 / 8000	8
NiCoTe ₂ //caC@pC	143 F·g ⁻¹ @ 1 A·g ⁻¹	50.84 Wh Kg ⁻¹	790.05W kg ⁻¹	87.88 /25000	9
Co ₂ Te ₂ @rGO//AC	64 mAh·g ⁻¹ @ 1 A·g ⁻¹	51 Wh kg ⁻¹	800 W kg ⁻¹	82 / 3000	11
NG-CoTe ₂ /CRF//AC	140.2 F g ⁻¹ @ 1 A g ⁻¹	49.8 Wh kg ⁻¹	985 W kg ⁻¹	125.2 / 5000	12
Cu _{0.5} Co _{0.5} Se ₂ // MXene	321 F·g ⁻¹ @ 1 A·g ⁻¹	84.19 Wh kg ⁻¹	715.12 W kg ⁻¹	91.1/10000	16
CoTe (4:1)//AC	169 F·g ⁻¹ @ 0.5 A·g ⁻¹	60 Wh kg ⁻¹	798 W kg ⁻¹	74 / 10000	17

CF@CoTe ₂ - NiTe ₂ //CF@Fe ₂ O ₃	131 F·g ⁻¹ @ 1 A·g ⁻¹	41 Wh kg ⁻¹	750 W kg ⁻¹	87.6 / 5000	18
1Td WTe ₂ Nanosheets - symmetric	221 F g ⁻¹	31 Wh kg ⁻¹	7.8 kW kg ⁻¹	91 / 5500	19
NiTe:Co//AC	103.4 F g ⁻¹ @ 1 A g ⁻¹	36.8 Wh kg ⁻¹	-	95 / 52000	20
1T'-MoTe ₂ /AC	158.9 F g ⁻¹ @ 1A g ⁻¹	56.4 Wh kg ⁻¹	0.8 k W kg ⁻¹	-	21
CoTe@C-NiF - symmetric	296.27 mF cm ⁻²	43.84 Wh kg ⁻¹	738.88 W kg ⁻¹	83.33 / 8000	22
CoOTe//AC	621 F·g ⁻¹ @ 1 A·g ⁻¹	72.82 Wh kg ⁻¹	0.635 kW kg ⁻¹	90.88/20000	23

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