Electronic Supplementary Material (ESI) for Journal of Materials Chemistry A. This journal is © The Royal Society of Chemistry 2023

Supporting information

$Ti_3C_2T_x$ and copper sulfide composite nanofluid with hierarchical structure for sustainable and efficient solar light-thermal conversion

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1. Experimental part

1.1 Light-thermal conversion performance test

In this work, the nanofluid was placed in a container, and illuminated by a solar simulator MICROSOLAR300 xenon light source (Beijing Perfectlight Technology Co., Ltd., China) with different light intensity, and its temperature changes were recorded through a data collector. Turn off the light source after the temperature reaches equilibrium, cool the nanofluid to room temperature. In addition, the temperature changes of nanofluids are recorded using an infrared thermal imager Fotric 225s.



Fig. S1 Light-thermal conversion performance testing device diagram.

1.2 Water evaporation performance test

The water evaporation experiment was performed using a solar simulator MICROSOLAR300 xenon light source (Beijing Perfectlight Technology Co., Ltd., China) outputting a simulated solar intensity of 1.0-3.0 kW/m² with an air mass (AM) 1.5 G filter and an interfacial evaporator as schematically illustrated in Fig. S2. Float the dialysis bag on the surface of 30 mL pure water, and 1.5 g of nanofluid was placed above the dialysis bag to form an interface evaporator. Finally, place the interface evaporator on an electronic balance (SY204, Shanghai Yoke Instrument Co., Ltd) with an accuracy of ±0.1 mg to detect changes in water quality. Use CEL-NP2000 optical power meter (China Education Au light Technology Co., Ltd., China) to detect changes in light intensity. The water evaporation rate (E) and solar-to-vapor conversion efficiency (η_E) are calculated using the following equation:

$$E = \frac{\Delta m}{S \cdot t} \tag{S1}$$

$$\eta_E = \frac{E \cdot H_{L-\nu}}{S_m} \tag{S2}$$

$$H_{L-v} = H_s + H_e \tag{S3}$$

$$H_s = C(T_i - T_0) \tag{S4}$$

where Δm is the mass loss of water, S is the illuminated area of the evaporator, t is the illumination time, S_m is the illumination intensity. In addition, H_{L-v} is the enthalpy of liquid-vapor phase transition. H_{L-v} was calculated by eq. S3 and S4, where H_s is the sensible heat per unit mass of water, C is the specific heat capacity of water (4.2 J/(g·k)), T_i is the final temperature of the sample surface, and T_0 is the initial temperature of water. H_e is the phase transition enthalpy of water at T_i .



Fig. S2 Schematic diagram of water evaporation performance testing device.

1.3 Preparation of [EBIM]NTf₂

[EBIM]NTf₂ was prepared according to the methods reported in previous literature. Firstly, 150 mL of ethylacetate was measured and place it in a three necked flask as the reaction solvent. Subsequently, 0.01 mol 1-butylimidazole and 0.012 mol bromoethane were added, and reacted 24 h at 55 °C after mixed evenly. After the reaction was completed, the liquid undergoes layering phenomenon after being allowed to stand for a period of time. [EBIM]Br was obtained through wash the lower sediment repeatedly with ethylacetate, and dried it in a 70 °C oven. Afterwards, [EBIM]NTf₂ was prepared through anion exchange. Specifically, 0.01 mol [EBIM]Br was dispersed in H₂O, and then added to 50 mL of H₂O containing 0.011 mol of LiNTf₂. The reaction was carried out under magnetic stirring at 65 °C for 24 h, and the resulting lower layer precipitate can be prepared by washing and drying with water.

2. Characterization of MXene



Fig. S3 Selected area electron diffraction diagram of FCuS.



Fig. S4 (a) SEM image of accordion shaped MXene. (b) AFM image of $Ti_3C_2T_x$ nanosheet. (c, d) TEM and SEM images of $Ti_3C_2T_x$. (e) The Tyndall effect of MXene



Fig. S5 (a) The XRD patterns of Ti_3AlC_2 and $Ti_3C_2T_x$. (b) The wide scan spectra of XPS of Ti_3AlC_2 and $Ti_3C_2T_x$ and elemental scan of (c) Ti, (d) C, (e) O and (f) F.

3. Characterization of [EBIM]NTf₂



Fig. S6 Schematic diagram of [EBIM]NTf₂ preparation.



Fig. S7 (a) ¹H NMR spectrum and (b) FT IR spectra of bromoethane, 1butylimidazole, [EBIM]Br and [EBIM]NTf₂.

4. Characterization of Ti₃C₂T_x/FCuS-IL and Ti₃C₂T_x/HCuS-IL nanofluids



Fig. S8 (a, b) TEM images of $Ti_3C_2T_x/FCuS-PDA$. (c) The mapping analysis results of $Ti_3C_2T_x/FCuS-PDA$. (d, e) TEM images of $Ti_3C_2T_x/HCuS-PDA$. (f) The mapping

analysis results of Ti₃C₂T_x/HCuS-PDA.



Fig. S9 The TGA curves of $Ti_3C_2T_x$, FCuS, HCuS, $Ti_3C_2T_x$ /FCuS, $Ti_3C_2T_x$ /HCuS and

PVP-K30.

Sample	Specific surface area (m ² /g)
$Ti_3C_2T_x$	11.677
FCuS	88.95
HCuS	28.226
Ti ₃ C ₂ T _x /FCuS	152.707
Ti ₃ C ₂ T _x /HCuS	120.682
$Ti_3C_2T_x$ /FCuS-PDA	238.319
Ti ₃ C ₂ T _x /HCuS-PDA	211.199

Table S1 Specific surface area of different samples



Fig. S10 Infrared thermal imaging of the cold platform surface.



Fig. S11 Optical photos of H_2O , [EBIM]NTf₂, and $Ti_3C_2T_x$ /FCuS-IL (0.12%)

nanofluid placed on a cold bench for 60 s.



Fig. S12 Optical photos of [EBIM]NTf_2 and $Ti_3C_2T_x$ /FCuS-IL (0.12%) nanofluids

dripping with a 5 mL syringe



Fig. S13 (a) The viscosity-temperature curves, (b) modulus-temperature curves and (c) modulus-angular frequency curves of $Ti_3C_2T_x/HCuS$ -IL. (d) The Zeta potential of $Ti_3C_2T_x/FCuS$ -IL and $Ti_3C_2T_x/HCuS$ -IL.



Fig. S14 Absorption of (a) Ti₃C₂T_x/FCuS-IL and (b) Ti₃C₂T_x/HCuS-IL, respectively.



Fig. S15 Temperature dependence curves of (a-c) $Ti_3C_2T_x/FCuS$ -IL and (d-f) $Ti_3C_2T_x/FCuS$ -IL under different light intensities irradiation (0.5 Sun, 2 Sun, 3 Sun).



Fig. S16 Infrared thermal imaging of [EBIM]NTf₂, $Ti_3C_2T_x$ /FCuS-IL and $Ti_3C_2T_x$ /HCuS-IL with different mass fraction of $Ti_3C_2T_x$ /FCuS and $Ti_3C_2T_x$ /HCuS after irradiated at different light intensities (0.5 Sun, 1 Sun, 2 Sun and 3 Sun) for 1 h.



Fig. S17 Temperature dependence curves of (a) FCuS-IL, (b) $Ti_3C_2T_x$ -IL and (c) $Ti_3C_2T_x$ /FCuS-IL under different light intensities irradiation. (d) T_{max} of CuS-IL,





Fig. S18 The cyclic η_{LTC} of (a) Ti₃C₂T_x/FCuS-IL (0.12%) and (b) Ti₃C₂T_x/HCuS-IL (0.12%) under different solar intensity (0.5 Sun, 1 Sun, 2 Sun and 3 Sun).



Fig. S19 Infrared thermal imaging changes of [EBIM]NTf₂ IL under different solar intensities for 5 min.



Fig. S20 Mass change of water within 60 min of exposure to different light intensities (a) 0.5 Sun, (b) 1 Sun, (c) 2 Sun and (d) 3 Sun.



Fig. S21 Separation and reused the photoabsorbers after the experiments

Table S2 Comparison of light-thermal conversion performance with previously

reported literature

Materials	Test conditions	ΔT (°C)	η (%)	References
MXene nanofluid	1 Sun 0.002%	-	63.35	1
MXene nanofluid	1 Sun 0.02%	22	80	2
AuNP solutions	1 Sun	6	80	3

	0.2 mM				
TiO ₂ nanofluid	1 Sun 2.0%	15	40	4	
Co@NC-900 nanofluid	1 Sun 0.01%	25	45	5	
Ti ₃ C ₂ T _x /PDA-IL	1 Sun 0.04%	47.9	83.93	6	
magnetic FCNTs nanofluids	2 Sun 0.01%	11.5	_	7	
TiN/MWCNTs hybrid nanofluids	1 Sun 0.004%	11	76.4	8	
ZrC/TiN nanofluids	1 Sun 0.016%	22	73.7	9	
Ag@Fe ₃ O ₄ nanofluids	2 Sun 0.03%	23	_	10	
MWCNT-DW/EG nanofluids	1 Sun 0.05%	38	92.0	11	
Fe ₃ O ₄ /CNTs nanofluids	2 Sun 0.015%	14	92.1	12	
PR nanosheets nanofluid	1 Sun 1.0%	50	-	13	
Ti ₃ C ₂ T _x /FCuS-IL	1 Sun 0.08%	60.2	94		
Ti ₃ C ₂ T _x /HCuS-IL	1 Sun 0.10%	56.3	88.1	I IIIS WOFK	

Table S3 Comparison of water evaporation performance with previously reported

literature				
Materials	Evaporation	Water	Solar-to-vapor	References

	conditions	evaporation rate	conversion		
		$(kg/(m^2 \cdot h))$	efficiency (%)		
PDA@MXene	1 Sun	1.276	85.2	14	
	3 Sun	4.0	90%		
carbon aerogels	1 Sun	1.29	87.51	15	
CDS T: C	1 Sun	1.32	91.9	16	
CDS-11 ₃ C ₂	3 Sun	3.84	88.2		
G1 MXene nanocoating	1 Sun	1.37	-	17	
MXene/PDA@	1 Sun	2.09	94.44		
TiO ₂ /Fe ₃ O ₄ @ C22-HMC	3 Sun	6.36	95.85	18	
Janus hydrogel- 1	1 Sun	1.45	_	19	
CB-Wood	1 Sun	2.52	75	20	
Janus PMX membrane	1 Sun	1.4	80	21	
PPy/MXene- PDA-fabric	1 Sun	1.55	90	22	
	1 Sun	1.5	91	23	
MXene/CIS	2 Sun	2.5	80	23	
MXene/cellulos e	1 Sun	1.44	91.5	24	
Fe ₃ O ₄ /CNTs nanofluids	2 Sun	0.9	89	12	
Ti ₃ C ₂ T _x /FCuS-	1 Sun	1.34	93		
IL (0.12%)	3 Sun	2.26	53.8	This work	
Ti ₃ C ₂ T _x /HCuS-	1 Sun	1.26	87		
IL (0.12%)	3 Sun	2.03	48		

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