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Solar-driven seawater desalination *via* plasmonic hybrid MOF/polymer and its antibacterial activity†

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In recent years, solar seawater desalination has been considered to be a promising and cost-effective technique to produce clean sources for water treatment and water deficiency. In addition, this technique shows high photothermal conversion efficiency by solar collectors to transfer solar energy into heat and the transformation of molecules in the capillaries of solar evaporators. In this study, we report the preparation of graphene-supported MIL-125 with polyurethane foam (MGPU) for solar steam generation. We modified MGPU by using the plasmonic nanoparticles of Ag and a polymer of polyaniline to increase the evaporation rate. Polyurethane foam can float on the surface of water and self-pump water by its hydrophilic porous structure, superior thermal insulation capabilities, and easy fabrication. MIL-125 has a high salt rejection and higher water permeability. It can reduce the affinity between water molecules and the pore surface of membrane, making it simple for water molecules to move through the pores. GO is a great alternative for steam generation applications since it exhibits broad-band light. The strong solar absorption, photothermal conversion efficiency, and photoreaction efficiency are enhanced by the use of silver nanoparticles in the photoreaction. The salt resistance capability is enhanced in saline water in the presence of polyaniline in a composite. Under one solar irradiation, the Ag/PANI/GO@MIL-125 (Ag-PMG) nanocomposite demonstrates an average $1.26 \text{ kg m}^{-2} \text{ h}^{-1}$ rate of evaporation and an efficiency as high as 90%. The composite exhibits remarkable stability and durability after more than 10 cycles of use without a noticeable decrease in activity. In addition, the composite exhibits excellent organic dye removal from contaminated water and generates pure condensed freshwater. The antibacterial photoactivity of the photocatalysts was examined against *B. subtilis* and *E. coli*. The results demonstrate that Ag-PMG shows higher antibacterial activity than MIL-125 and PMG. It was shown that the presence of rGO, PANI, and Ag in the sample enhances the antimicrobial activity.

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1. Introduction

Freshwater supplies are currently being depleted due to high wastewater discharge and industry usage.¹ Currently, the decrease in pure water is one of the important global problems that threaten humanity and needs to be solved.² Pure water is necessary for different uses, including drinking, agriculture, and industrial uses.³ According to the United Nations Environmental Program (UNEP), one-third of the world's population lives in water-stressed areas, and two-thirds of the global population will endure water scarcity by 2025.⁴ The population and energy needs will continue to rise, accompanied by economic expansion. By 2035, global energy consumption is expected to rise by 31%.⁵ Desalination is one technique for extracting pure water from salty, brackish, or contaminated water. All

desalination techniques can be used to resolve the freshwater source shortages. These processes, which include multi-effect desalination (MED), reverse osmosis (RO), desalination by humidification and dehumidification (HD), multi-stage flash distillation (MSF), membrane distillation (MD), mechanical vapour compression (MVC) desalination, and low-temperature desalination (LTD), frequently involve high energy consumption, high material requirements, and high costs.^{6–8} We can improve the evaporation efficiency and use alternative energy sources for heating in seawater desalination, such as solar energy.^{9,10} The most renewable energy source is solar energy, which has large technical feasible potential (about 60 TW) among all sources of renewable energy. The solar steam generation strategies can easily transform light into heat water with low photothermal conversion efficiency because of the significant heat loss.¹¹ Three categories of using solar energy are currently available, such as photothermal conversion (PTC), photoelectric conversion,¹² and photochemical conversion.¹³ For heat insulation and liquid supply, a substrate and high-efficiency PTC materials are required.¹⁴

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Photothermal conversion materials (PCMs) are cost-effective and simple methods to generate steam that floats at the water–air interface, which may produce heat from solar energy. Additionally, the PCMs can produce solar steam and freshwater, localize heat in the upper region, and decrease heat loss to the bulk water. It is necessary to produce photothermal conversion materials with small thermal conductivity, self-floating properties and low cost.^{15–22} Light absorption by the photothermal layer is important for large-efficiency solar desalination. Plasmonic metals for surface plasmon resonance (SPR), such as gold,²³ silver,²⁴ and alumina,²⁵ can be used for efficiently heating water by increasing the temperatures over its boiling point, and can absorb more than 95% of the sunlight for solar desalination. We demonstrate other solar absorbers with surface plasmon resonance to enhance the rate of evaporation with metal–organic frameworks, graphene oxide and polymer.

Metal–organic frameworks (MOFs) are important candidates because of their high porosity, charged surface, hydrophilicity and evaporation-driven electricity generation, such as MIL-125 for efficient water desalination. Titanium-based MOFs have high porosity, structural stability, and high electrical and optical properties, nontoxicity, and are low-cost. Their photocatalytic applications include H₂ production, CO₂ reduction, H₂O₂ production, organic and inorganic waste remediation, and N₂ fixation.^{26–30} Lately, the use of metal–organic frameworks (MOFs) as new materials during water desalination has attracted much attention among engineers. MIL-125(Ti) has light response characteristics due to the development of new Ti-MIL structures and their application in many fields.³¹ The organic ligands in MOFs can be stimulated by light and transport electrons to the metal center, making them useful as semiconductor-like photocatalysts.³² MIL-125 has excellent efficiency, water permeability and salt rejection in the desalination process.^{33–36} MIL-125 and its modification can be used in photocatalytic processes.^{37–45} Titanium MOFs have important research applications as anti-inflammatory, antibacterial and anticancer agents.

For solar desalination, graphene oxide (GO) has also been described as a photothermal material due to its effective light absorption, highly controlled microstructures, and production simplicity. GO is a great candidate for steam generation systems, including optoelectronic and biological applications, since it exhibits a broad spectrum of light absorption from the visible to NIR region. By the size effect, graphene oxide (GO) helps to reject hydrated salt ions.^{46,47} Graphene oxide (GO) and rGO are necessary for toxicity to Gram-positive and Gram-negative bacteria.⁴⁶ Polyaniline (PANI) is known for its flexibility, high conductivity, good light absorption property, chemical stability, and solar-heating interfacial evaporation.⁴⁹ PANI is used in many fields, including electrostatic discharge systems, batteries, and chemical and biochemical sensors. PANI has high antibacterial activity because of the electrostatic interaction between PANI and bacteria, and the increase in the rate of the host materials for catalysis and antibacterial activity.^{50,51} The Ag and PANI materials have shown antibacterial activity against *B. subtilis* bacteria, and has also been reported against another bacteria.⁵²

Herein, we report the fabrication of nanostructured MIL-125 as solar absorbers because of its large solar absorptance, and cost-effective synthesis. To enhance the evaporation performance, we report a solar thermal evaporator composite of plasmonic GO-based MIL-125. The conductivity, thermal stability and light absorption property were enhanced by modification using PANI and Ag. In this paper, we prepared a floating polyurethane foam (PU), which is mixed with a photothermal composite by a simple method.

2. Experimental

2.1. Materials and chemicals

All solvents and chemicals were supplied and used without purification. Titanium(IV) isopropoxide (TTIP), 1,4-benzene dicarboxylic acid (H₂BDC), *N,N*-dimethylformamide (DMF) and methanol were acquired from Sigma Aldrich. For the synthesis of GO, the following materials were used as follows: graphite powder (99.95%), potassium permanganate (KMnO₄), sodium nitrate (NaNO₃), sulfuric acid (H₂SO₄), and 30% hydrogen peroxide (H₂O₂). Hydrochloric acid (HCl), absolute ethanol silver nitrate (AgNO₃), sodium borohydride, ammonium persulfate and aniline were used for surface modification.

2.2. Synthesis of graphene oxide (GO)

Graphite flakes were used to prepare graphene oxide by using a modified Hummer's technique. In a 250 mL beaker, 115 mL of concentrated H₂SO₄ was added at 0 °C (in an ice bath). A mass of 2.5 g ground NaNO₃ was added to the previous solution, then 4.0 g graphite powder was added after all of the ground NaNO₃ was completely dissolved. After a duration of 20 min, 15.0 g KMnO₄ was added to the mixture very slowly, ensuring that the temperature did not exceed 5 °C. After 15 min, the heat of the mixture reached 35 °C in 3 h and transferred to 2 L beaker, followed by the addition of 230 mL DI H₂O. A volume of 700 mL DI H₂O was added after 20 min at a temperature of 80 °C. A volume of 20 mL 10% H₂O₂ was poured into the mixture after 20 min, and the color changed to golden yellow. The resulting graphene oxide was washed with 2 L acidified water (HNO₃), followed by 5 L hot DI H₂O (80–100 °C) at 1 L each time. The brown emulsion was put into a convection oven at 60 °C for drying.

2.3. Synthesis of MIL-125

By using the solvothermal method, we synthesized MIL-125 depending on the following procedures. 1,4-Benzene dicarboxylic acid (15 mmol) was dissolved in a mixture of methanol (5 mL) and anhydrous DMF (45 mL). Subsequently, TTIP (9 mmol) was added to the mixture and stirred for 10 min. The mixture was placed into a Teflon tube and heated at 120 °C for 16 hours, and put in a conventional oven. The autoclave was cooled to reach room temperature. Finally, we collected the white MIL-125 by using a centrifuge, and washed the precipitate several times with DMF and then diluted ethanol. Finally, the white solid was dried at 80 °C, and maintained in a small glass vial.



RSC Adv., 2023, 13, 18525-18537 | 18527

$$\eta = \frac{H_e v}{Q_{in}} \quad (2)$$

which n is the rate of vapor formation during steady-state conditions irradiation ($\text{kg m}^{-2} \text{h}^{-1}$), and Q denotes the intensity of the incoming solar irradiation (mW cm^{-2}). H_e is the result of adding the sensible and latent heat enthalpies, which is calculated as follows:

$$H_e = C_p \Delta T + \Delta H \quad (3)$$

where, C_p is water's specific heat capacity ($4.18 \text{ J g}^{-1} \text{K}^{-1}$), ΔT is water's temperature change, and ΔH is the latent heat of vaporization (2394 kJ kg^{-1}) on the relative temperature.

2.11. Water solar desalination artificial seawater and wastewater test

The preparation of the artificial seawater sample, which contains NaCl (1000 ppm), $\text{MgSO}_4 \cdot 7\text{H}_2\text{O}$ (1500 ppm), and CaCl_2 (1000 ppm), involved dissolving deionized water in saltwater that had undergone solar desalination. By using data from inductively coupled plasma optical emission spectroscopy, we showed the concentrations of the principal ions (Na^+ , Mg^{2+} , and Ca^{2+}) in the saltwater. In order to obtain polluted wastewater, 10 ppm of methylene blue and rhodamine B were also prepared.

2.12. Solar steam generation recycling test

Ten solar steam generation cycles were performed to assess the reusability of the Ag-PMG nanocomposite foam. Using the solar simulator system, the samples were exposed to 1 kW m^{-2} of light for each cycle. The values of water mass loss and the surface temperature were calculated after one hour in 1 sun. The wetted polyurethane foam was cleaned, dried, and then reused for the following cycle.

3. The antibacterial activity

The antibacterial activities of the fabricated samples were tested against Gram-positive *B. subtilis* and Gram-negative *E. coli* bacteria by agar disc diffusion technique using inoculums containing 106 bacterial spread on Muller–Hinton agar plates. The sample discs were placed on the surface of the agar plates seeded with the test organisms. The plates were incubated at 37°C . Diameters of inhibition zones (mm) were measured after 18–24 hours for bacteria.

4. Result and discussion

4.1. Characterization

Fig. 1b shows the XRD patterns of the synthetic composites and pure MIL-125. The XRD pattern of pure MIL-125 has prominent peaks that were in excellent agreement with our previously published literature, and demonstrate the good crystallinity of the frameworks.⁵⁴ The MIL structure with 3D pores was created and connected to the sharing of $\mu\text{-OH}$ corners with TiO_6 octahedral interconnected to linker (H_2BDC) molecules. For pure MIL-125, the peaks allocated at $2\theta = 6.9^\circ, 9.9^\circ, 11.9^\circ, 15.7^\circ, 18^\circ,$

$19.2^\circ, 22.8^\circ, 23.7^\circ, 34.3^\circ$ and 49.3° are assigned to the (101), (002), (211), (220), (310), (222), (312), (213), (400) crystal planes, respectively.^{55–58} This result demonstrated that MIL-125 crystals were completely formed under the influence of GO sheets.⁵⁹ The XRD pattern of the presence of polyaniline in the composite illustrated the same crystal structure of MIL-125 during the fabrication of the composite.⁶⁰ There are apparent weak diffraction peaks at 38.1° and 44.2° in the Ag-MG composite, indicating that Ag is loaded on the composite's surface,⁶¹ so the Ag peaks were magnified and are labeled in Fig. S1† to show the diffraction peak.

The functional groups of all fabricated composites were observed by using FTIR spectra, as depicted in Fig. 1a. The significant broad adsorption band in the $3200\text{--}3400 \text{ cm}^{-1}$ range was shown in the FTIR spectra of MIL-125 for free solvent molecules in the framework pores.⁶² The bands at $1291\text{--}1690 \text{ cm}^{-1}$ were produced by the vibration of carboxylate groups due to the linker in the frameworks.⁶³ The characteristic absorption band of the benzene ring was shown in the $800\text{--}1200 \text{ cm}^{-1}$ range, while the bands at $500\text{--}800 \text{ cm}^{-1}$ belonged to Ti–O–C and Ti–O–Ti vibrations.⁶⁴ The absorption band increased around 3400 cm^{-1} and 1600 cm^{-1} due to the O–H vibration and Ti–O–C bonds, respectively, owing to the interactions between MIL-125 and GO nanosheets.⁶⁵ A new absorption band around 1000 cm^{-1} was assigned as the $\pi\text{--}\pi$ interaction between the frameworks and GO sheets. In addition, the absorption band originating from the graphene skeletal vibration was observed at 1626 cm^{-1} . New bands at 1125 and 805 cm^{-1} belonged to the C–N stretching and C–H out-of-plane bonding in the benzenoid ring, respectively.^{66,67} In addition, an absorption band was observed at 3251 cm^{-1} due to the amine group stretching.^{68,69} The band intensity decreased compared to MG due to $\pi\text{--}\pi$ interaction between PANI with graphene oxide, and hydrogen bonding between them. The peak at 1627 cm^{-1} exhibited low intensity compared to MG due to AgNPs on the graphene oxide surface.

The elemental states and chemical compositions of all composites have been estimated using HR-XPS analysis. Fig. 2 shows the XPS of the Ag-PMG composite that determined the existence of C 1s, O 1s, N 1s, Ti 2p and Ag 3d elements. HR-XPS of the C 1s pattern was deconvoluted into five peaks at 284.69, 285.92, 287.21, 288.92 and 290.83 eV, attributed to C–C (sp^2 , sp^3), C–N, C=O, and –COOR, respectively.^{70–72} The O 1s spectrum was divided into three main peaks at 530.28, 531.97, 532.96 and 535.93 eV that were ascribed to metal–oxygen bonds (O–Ti–O), –OH in the organic ligand and oxygen adsorbed at the surface (C=O) and H_2O molecules, respectively.⁷³ The N 1s spectrum was divided into three peaks at 401.36, 399.64 and 398.80 eV binding energies, which belonged to the (N^+), quinoid imine bonds (–NH–) and benzenoid amine bonds (=N–) of polyaniline, respectively.^{74,75} The peaks at 459.09 and 464.75 eV resulted from the deconvolution of the Ti 2p spectrum to two peaks belonging to Ti $2p_{1/2}$ and Ti $2p_{3/2}$, respectively.⁷⁶ The Ag 3d spectrum exhibited two peaks assigned to Ag $3d_{3/2}$ and Ag $3d_{5/2}$, and the spin-orbital photoelectron peaks are shown at 367.20 and 373.23 eV, respectively.^{77,78}



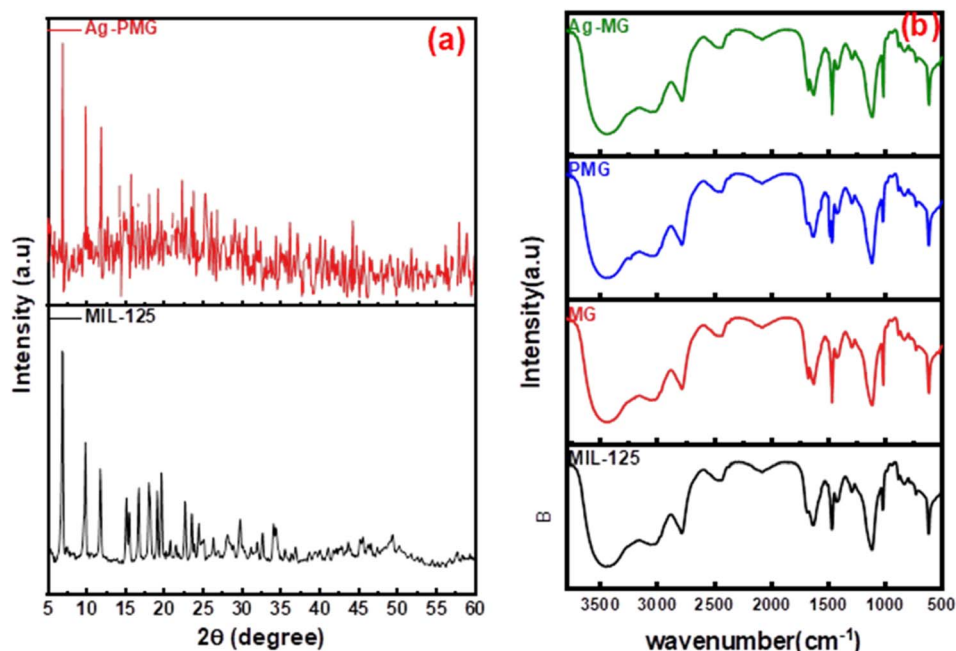


Fig. 1 (a) XRD patterns and (b) FTIR spectra of pure and modified MIL-125.

The morphology of the prepared composite was studied by SEM analysis before and after loading on the PU foam. Fig. 3 displays the structural information of the Ag-PMG composite, as determined by SEM. The high porosity of the polyurethane foam is shown in the SEM images, which contain the open porous, hydrophilic nature and cellular morphology of the PU matrix. The characteristics of the PU foam are important for water supply from the porous surface with low heat loss to bulk

water for solar evaporation.^{79–81} MIL-125 features a spherical shape with a smooth surface. From the SEM images, it can be seen that GO and PANI have a sheet-like shape. PANI was uniformly dispersed throughout MG and exhibited a deep layer with a pore structure, as well as numerous fold structures and agglomerations on the layer's surface. With the addition of Ag NPs, the surface of the material becomes rough and randomly deposited on the surface of PMG.^{82–85} Moreover, the surface area

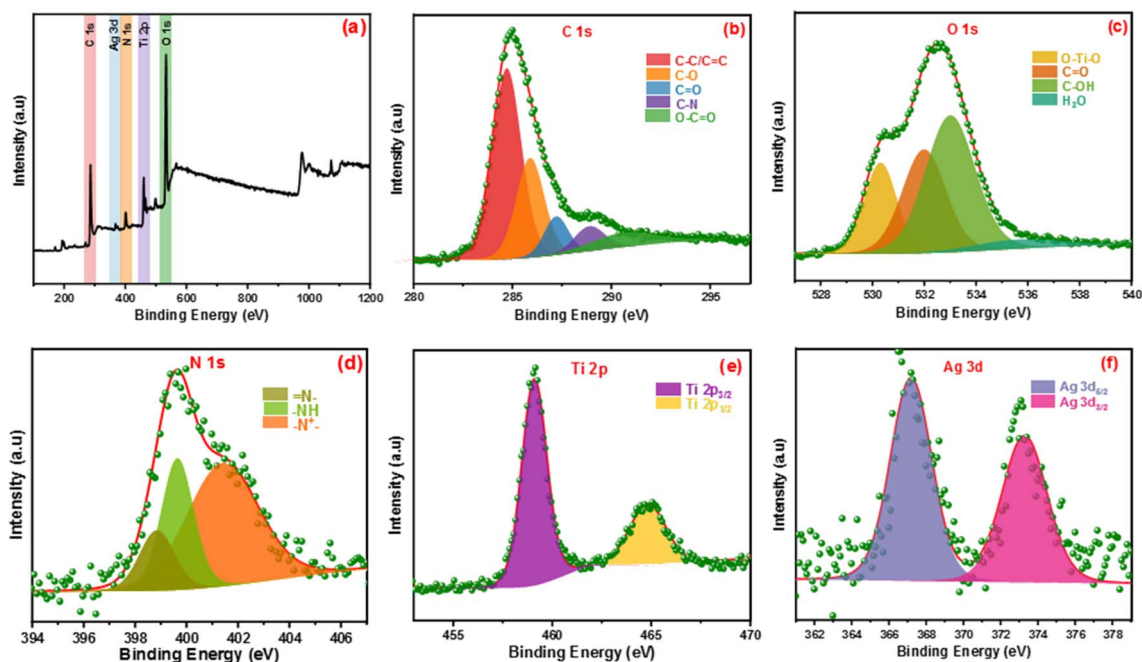


Fig. 2 (a) Full survey XPS spectrum of Ag-PMG, (b–f) high-resolution XPS of C 1s, O 1s, N 1s, Ag 3d and Ti 2p, respectively.

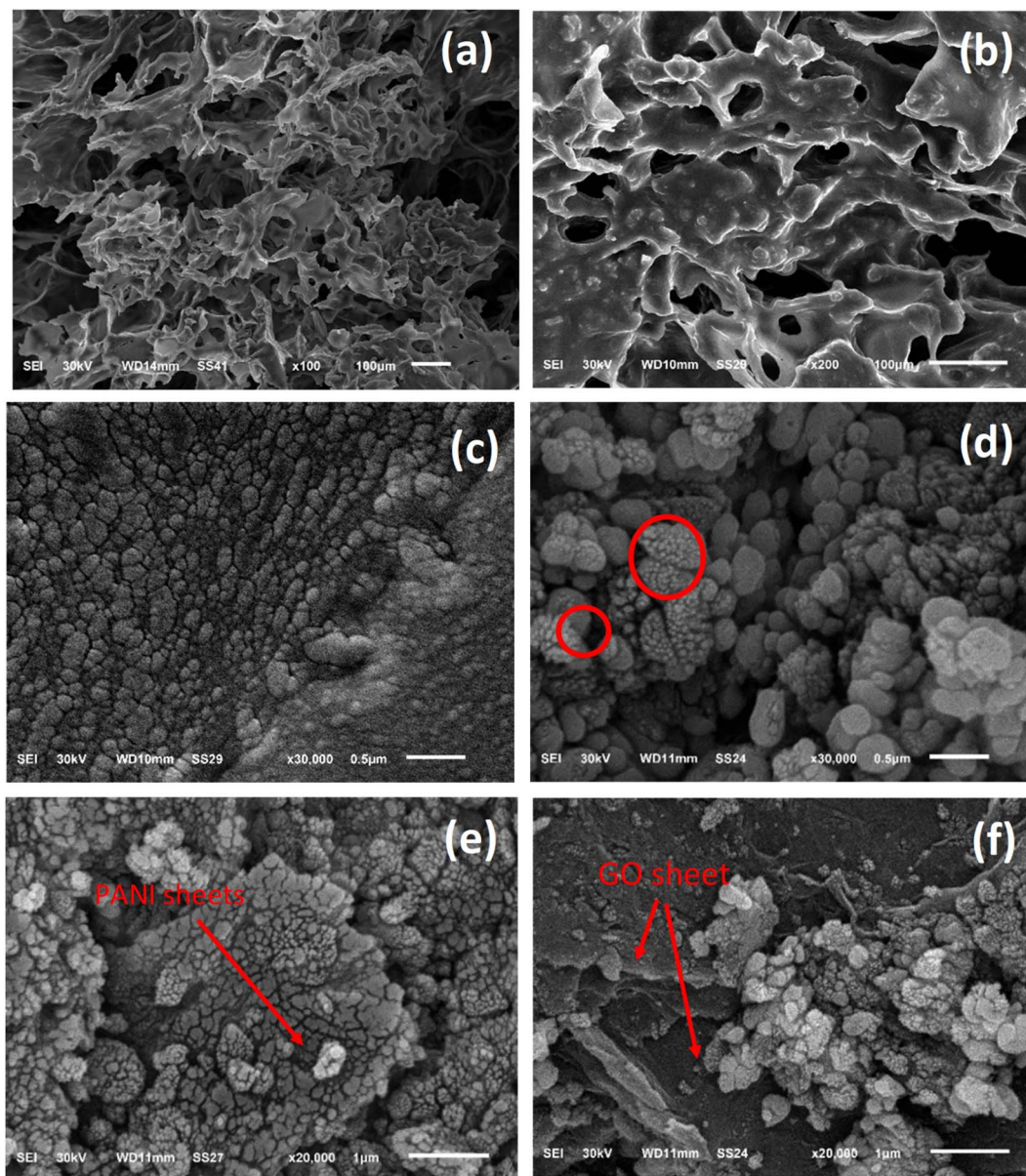


Fig. 3 SEM images of (a) PU foam, (b and c) Ag-PMG loaded on PU foam, (d–f) Ag-PMG composite.

was measured for pristine MIL-125, which shows a typical type (I) curve with a high surface area of $1120 \text{ m}^2 \text{ g}^{-1}$. Loading the sample with PANI and Ag nanoparticles decreased the surface area to $560 \text{ m}^2 \text{ g}^{-1}$ without affecting the porosity of the prepared material (Fig. S2†).

4.2. Solar steam generation process

To determine the steam generation performance for various fabricated photothermal absorbers, five polyurethane foams with a diameter of 4.5 cm were made using the following samples: M, MG, Ag-MG, PMG and Ag-PMG. All photothermal absorbers were demonstrated and contrasted by using polyurethane foam in 1 sun over 1 h by the solar simulator for the solar steam generation performance. Each sample was mixed with polyurethane foam with 4.5 cm inner diameter. The foam

was moistened on the surface of a volume of 100 mL water in a beaker, and all were wetted in deionized water before putting on the water surface. The mass loss was calculated by using an electronic balance, and the evaporation rate was measured from the mass loss was determined at 5 min up to 1 h.

The mass loss of five photothermal absorbers of the pure water (polyurethane foam) from the steam generation test is depicted in Fig. 4a. The results show that pure PU foam has the smallest mass loss of water of 0.31 kg m^{-2} between another sample. This is because pure PU foam has a low thermal absorptivity, but also moved the light absorption by its modification on the surface foam.^{86,87} The mass change of the (M, MG, PMG, Ag-MG and Ag-PMG) absorbers are 0.44, 0.26, 0.85, 0.97 and 1.26 kg m^{-2} , respectively, which demonstrated the combination of the frameworks with GO and plasmonic



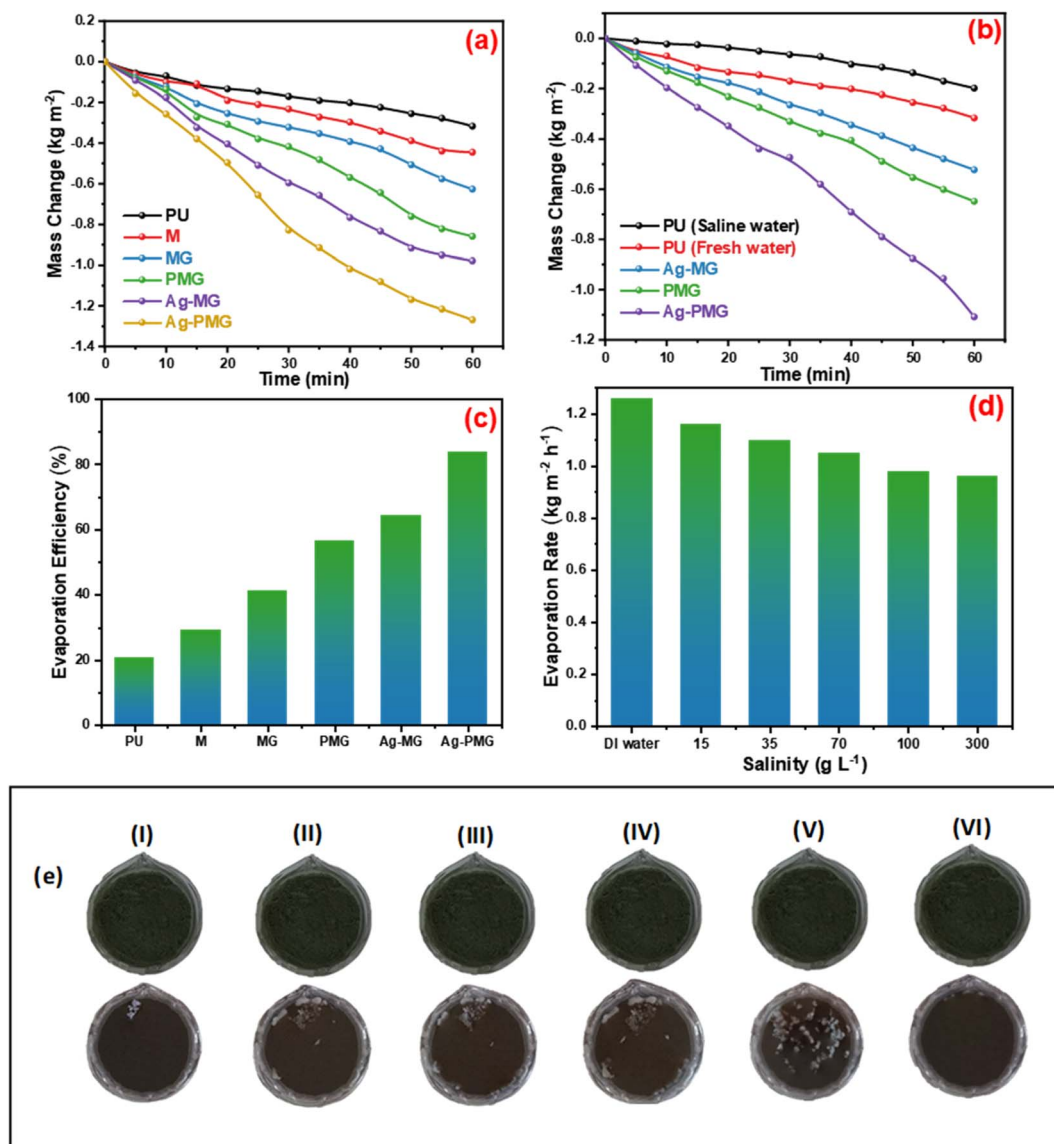


Fig. 4 (a) Mass changes for PU, M, MG, PMG, Ag-MG and Ag-PMG. (b) Mass change for Ag-MG, PMG and Ag-PMG of pure and saline water. (c) The evaporation efficiency of fabricated foams. (d) The evaporation rate of Ag-PMG for various concentrations of saline for 60 min, and (e) salt-rejection performance during a daytime salt aggregation time from (I) 5 h to (V) 9 h, and (VI) light-off salt diffusion times.

nanoparticles. MIL-125 was known as a photosensitive material and has a three-dimensional pore structure from phthalic acid and Ti-O octahedron. MIL-125 is a porous material with highly ordered crystal structures from organic linkers and metal ions with large active sites, which lead to excellent photocatalytic efficiency, water permeability and salt rejection in the desalination process. MIL-125 has good electron-hole separation performance because of its Ti-O clusters with isolated titanium oxide quantum dots.^{88,89} The solar absorbers including GO enhanced the rate of evaporation because of transmittance loss, the low thermal conductivity, and low reflectance of GO.⁹⁰ The hydroxyl and carboxylic functional groups in the graphene oxide structure were important for enhancing the evaporating water because the strong π - π interactions between the MOF and GO can effectively increase the electron-hole pair separation and

enhance the photocatalytic performance. The solar absorbers containing PANI increased the evaporation rate and the mass change due to good light absorption properties, chemical stability and high conductivity for solar-heating interfacial evaporation, unique photothermal conversion capability and good adhesive properties.^{91,92} The solar absorbers containing silver nanoparticles in Ag-MG and Ag-PMG increased the rate of evaporation in contrast to other absorbers because of their efficient plasmonic activity in the visible light region and broad light absorption band by the plasmonic nanoparticles. After the sample was prepared, the results demonstrated that the Ag-PMG composite has a high mass loss of 1.26 kg m^{-2} because of the incorporation of MIL-125, GO, PANI and Ag NPs. Fig. 4b shows the results of photothermal foams in the artificial saline water for PU, PMG, Ag-MG and Ag-PMG evaporators. The mass

The quality of the evaporators was assessed in Fig. 7a when artificial seawater was present, and a condensing unit was used to collect the evaporated water. ICP-OES was used to measure

Evaporator	Support	Solar intensity (kW cm ⁻²)	Evaporation rate (kg m ⁻² h ⁻¹)	Efficiency (%)	Reference
Ag/Au-GO	PU	1	1.00	63.0	79
Fe ₂ O ₃ -rGO	—	1	1.12	70.0	99
Modified graphene aerogel	—	1	1.2	76.9	100
Ag@TiO ₂ nanoparticles	A flexible filter membrane	1	0.9	68.6	101
C-Au-TiO ₂	Glass or polished Cu	1	0.97	68	102
rGO polyurathene nanocomposite foam	PU	1	1	65	103
Ag ₂ S@PANI/carbonaceous porous material	PU	1	1.1	91.5	104
NiCo _x S _y -PANI@GF	Glass fiber membrane	1	1.3	78.7	105
Acid-doped polyaniline	Polystyrene foam	1	1.3	80.7	106
rGO/TiO ₂ composite	—	1	1.2	—	107
Ag/PANI/GO@MIL-125	PU	1	1.26	90	This work

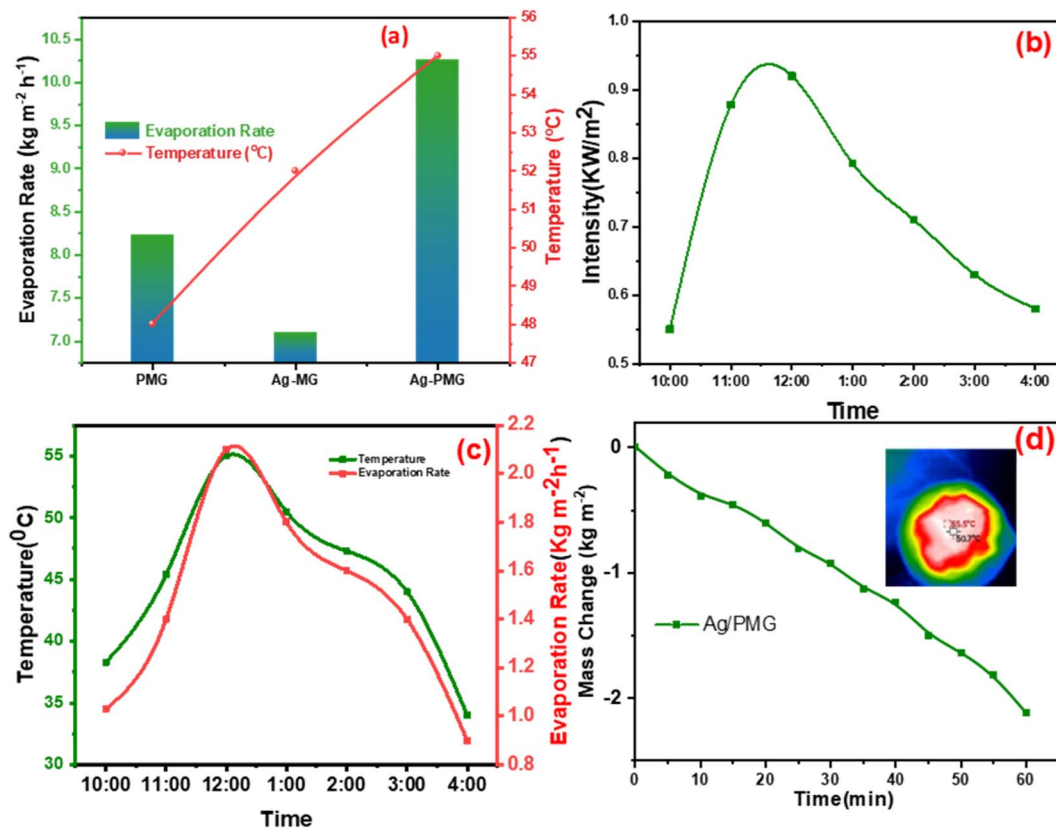


Fig. 5 Outdoor solar water evaporation system in real sunlight. (a) Evaporation rate during 7 h of outdoor solar desalination recorded over time from 10:00 to 4:00, and temperature of PMG, Ag-MG and Ag-PMG at high intensities. (b) Solar radiation intensities measured over time from 10:00 to 4:00. (c) Evaporation rate and temperature of Ag-PMG recorded over time from 10:00 to 4:00. (d) The mass change of Ag-PMG under the real sun for 60 min, and IR images of Ag-PMG at a high intensity of outdoor solar desalination.

the concentrations of Mg^{2+} , Ca^{2+} and Na^{+} ions in the condensed water. The concentration of the three ions decreased significantly after desalination with the Ag-PMG evaporator. The ion

concentrations decreased dramatically from their first levels of 1500, 1000, and 1000 mg L^{-1} of Mg^{2+} , Ca^{2+} and Na^{+} , respectively, which are below the standard values set by the World

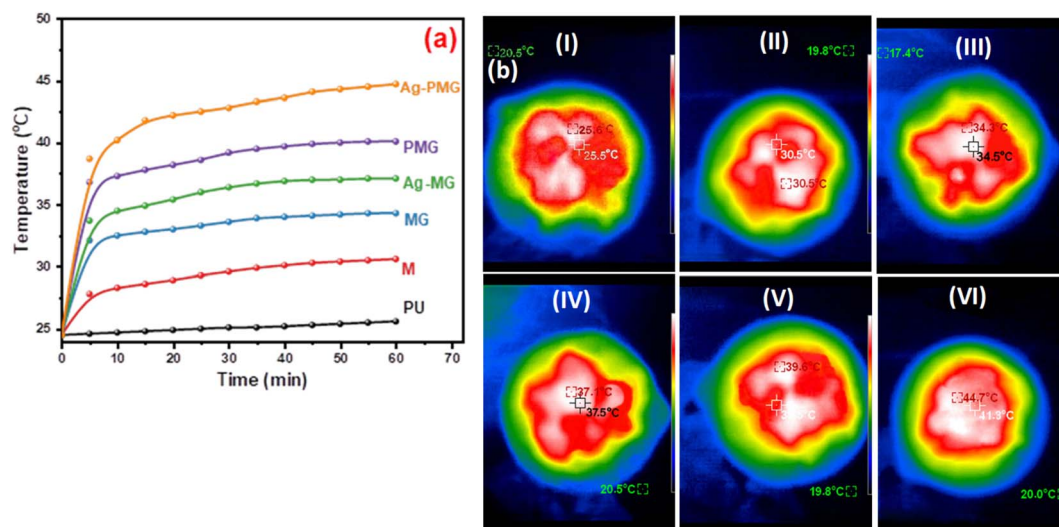


Fig. 6 (a) Temperature change on foam surfaces for different composites in 1 hour, and (b) IR thermal images of (I) PU, (II) M, (III) MG, (IV) PMG, (V) Ag-MG, and (VI) Ag-PMG after 1 h.

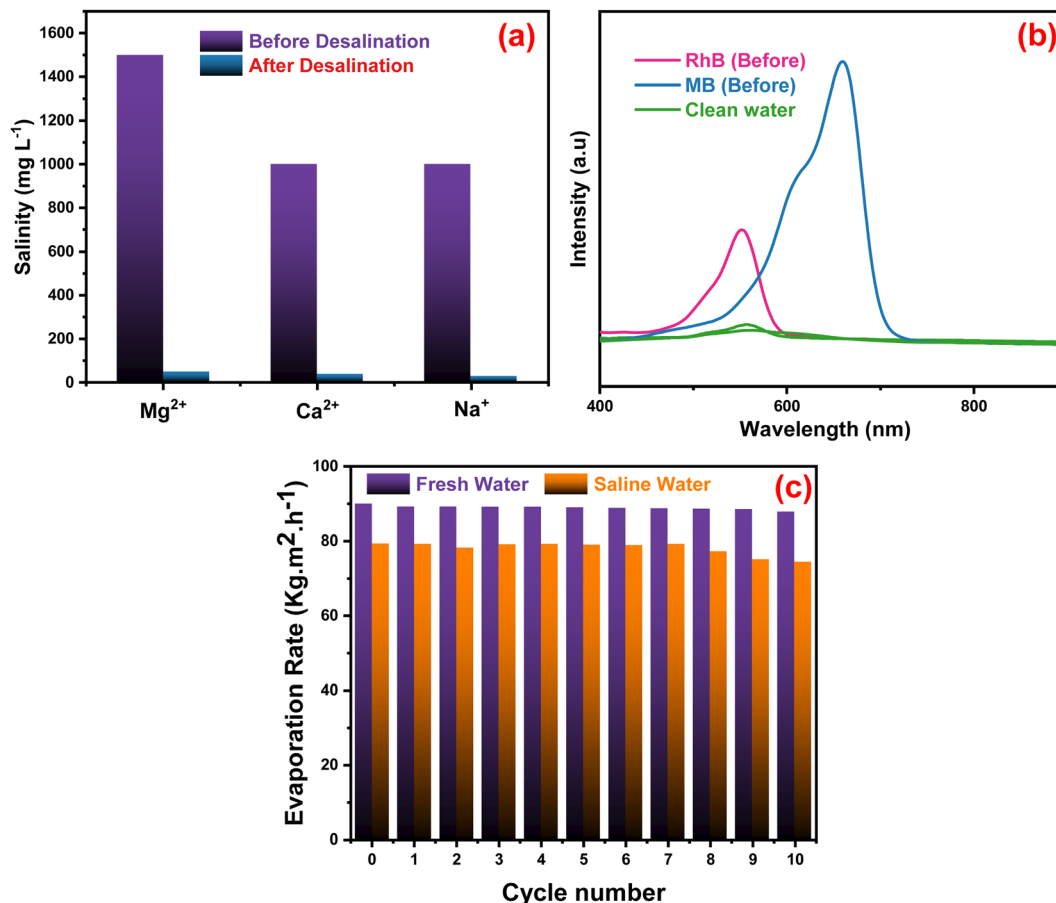


Fig. 7 (a) Concentration of metal ions before and after the desalination technique by using Ag-PMG composite, (b) UV-vis absorption spectra of dye-contaminated water before and after desalination, and (c) reusability test of the Ag-PMG sample during 10 cycles.

Health Organization for drinking water.¹⁰⁸ The synthesized system was investigated using excellent desalination techniques compared to common techniques. The Ag-PMG composite can be used to purify the contaminated water with RhB and MB dyes. Fig. 7b shows the absorption spectra of the polluted water before and after desalination, and demonstrated that all dyes have been eliminated from the evaporated water. Fig. 7c shows the reusability and stability of the Ag-PMG photoabsorber in seawater desalination by using 10 cycles of evaporated experiments in artificial seawater (1 hours of radiation for each cycle). The high efficiency of Ag-PMG after 10 cycles with a slight decrease can be attributed to plasmonic Ag nanoparticles and salt rejection of PANI, which prevent the blocking of the active sites of the solar absorber.²⁴

5. M, PMG and Ag-PMG composite tests as antibacterial agents

Antibacterial activity tests of M, PMG and Ag-PMG composites against both Gram-positive *B. subtilis* and Gram-negative *E. coli* bacteria were performed, and are shown in Fig. 8 and Table 2. MIL-125 has activity against *B. subtilis* but not against *E. coli* bacteria because the Gram-positive type gives more negatively

charged surfaces than the Gram-negative type, and MIL-125 has antibacterial activity against *B. subtilis* by producing a concentration of Ti^{4+} as an agent to produce positive charges. Its positive charges of Ti react with the negative charge cell membrane, affecting the structure of the cell membrane and changes its permeability, which leads to the death of the bacteria. GO has antibacterial activities due to the hydrophobic interaction with the membrane phospholipids. Ag-PMG composites have the highest antibacterial activity against both Gram-positive and Gram-negative bacteria by the inhibition

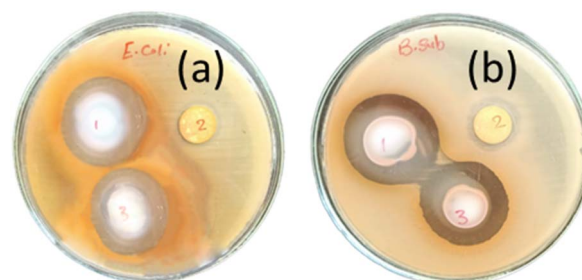


Fig. 8 Antibacterial activity tests of (1) Ag-PMG, (2) M, (3) PMG composites against both (a) *E. coli* and (b) *B. subtilis* bacteria.



Table 2 Antibacterial activity of M, PMG and Ag-PMG nanocomposites

Sample	<i>B. subtilis</i>	<i>E. coli</i>
	Inhibition zone (mm)	Inhibition zone (mm)
M	15	—ve
PMG	31.6	28.1
Ag-PMG	33.3	30.3

zone compared to the PMG and M composites because of the electrostatic adsorption between Ag and PANI molecules and bacteria cells, and the presence of silver nanoparticles kills bacteria. Ag reacts with the cytoplasmic membranes of the bacteria, leading to bacterial inactivation and death. Ag's spherical particles can easily enter cells through membrane holes, and the composite may shatter inside the cell due to reactions with sulphur and phosphorous-containing DNA and protein contents. The presence of silver ions bound to bacterial DNA disturbs bacterial respiration and adenosine triphosphate synthesis. Polyaniline and Ag have powerful antibacterial effects on bacterial strains. Also, it demonstrated stronger antibacterial defenses against the Gram-negative cell membrane by lowering the osmotic pressure, which supports their physical connection. Moreover, the increased permeability of the bacterial cell wall and the formation of stress-resistant biofilms are caused by the numerous chemical properties of polyaniline because of the surface hydrophilicity and the polymer chain length. The charged functional groups in the composite have an increased reaction with the bacteria cell. The inhibition zones of the studied bacteria are in the following order: *B. subtilis* > *E. coli*. This represents more activity of Ag and PANI on *B. subtilis* than on *E. coli*.^{48,109–112}

6. Conclusion

In summary, we prepared a new system of graphene/MIL-125 polyurethane (MGPU) nanocomposite foam for solar water desalination. The system has low thermal conductivity, porous structure, high absorption and the capacity for self-floating. MIL-125 has a pore structure, large active sites, water permeability, high surface area, and salt rejection in the desalination process. We can modify the MGPU foam by using plasmonic Ag and polyaniline. PANI improved the salt rejection capability and Ag improved the solar photothermal energy conversion in water desalination. The solar conversion efficiency of the Ag-PMG composite reaches 90% during 1 sun irradiation, and the evaporation rate is $1.26 \text{ kg m}^{-2} \text{ h}^{-1}$ connected to the physical properties of MIL-125 and GO, PANI and Ag. As a result, for continuous evaporation of 10 h, the evaporator showed stable evaporation. We showed the difference between the two composites, including PANI and Ag, and the effect of salt rejection in the composite. We also noticed that the composite has good durability after more than 10 cycles with no significant reduction in performance activity. The fabricated composites

can eliminate dyes from contaminated water and produce clean condensed freshwater. The presence of rGO, PANI and Ag in the fabricated catalysts enhances the antimicrobial activity.

Conflicts of interest

The authors declare no conflict of interest.

References

- Y. Xu, J. Ma, D. Liu, H. Xu, F. Cui and W. Wang, *Chem. Eng. J.*, 2019, **356**, 869–876.
- M. M. Mekonnen and A. Y. Hoekstra, *Sci. Adv.*, 2016, 1–6.
- K. H. Nayi and K. V. Modi, *Renewable Sustainable Energy Rev.*, 2018, **81**, 136–148.
- Z. Fang, Y. R. Zhen, O. Neumann, A. Polman, F. J. Garcia de Abajo, P. Nordlander and N. J. Halas, *Nano Lett.*, 2013, **13**, 1736–1742.
- K. Ramalingam, M. Liang, N. L. W. Pyae, S. H. Aung, T. Z. Oo, P. Srimuk, J. Ma, V. Presser, F. Chen and T. D. Waite, *ACS Appl. Mater. Interfaces*, 2020, **12**, 32788–32796.
- J. Lv, Z.-L. Chen, J. Tang, L. Chen, W.-J. Xie, M.-X. Sun and X.-J. Huang, *Surf. Coat. Technol.*, 2022, **446**, 128770.
- P. Srimuk, X. Su, J. Yoon, D. Aurbach and V. Presser, *Nat. Rev. Mater.*, 2020, **5**, 517–538.
- M. E. Suss and V. Presser, *Joule*, 2018, **2**, 10–15.
- M. Alduraibi, M. Hezam, B. Al-Ruhaimi, A. M. El-Toni, A. Algarni, M. Abdel-Rahman, W. Qing and A. Aldwayan, *Nanomaterials*, 2020, **10**(3), 413.
- M. Wang, J. Zhang, P. Wang, C. Li, X. Xu and Y. Jin, *Nano Res.*, 2018, **11**, 3854–3863.
- P. Tao, G. Ni, C. Song, W. Shang, J. Wu, J. Zhu, G. Chen and T. Deng, *Nat. Energy*, 2018, **3**, 1031–1041.
- X. Yang, W. Liu and P. Ren, *Phys. E*, 2016, **83**, 322–328.
- P. Liu, Y. Zhao, R. Qin, S. Mo, G. Chen, L. Gu, D. M. Chevrier, P. Zhang, Q. Guo, D. Zang, B. Wu, G. Fu and N. Zheng, *Science*, 2016, **352**, 797–801.
- M. H. Ahmadi, M. Ghazvini, M. Sadeghzadeh, M. Alhuyi Nazari, R. Kumar, A. Naeimi and T. Ming, *Energy Sci. Eng.*, 2018, **6**, 340–361.
- D. Weng, F. Xu, X. Li, Y. Li and J. Sun, *J. Mater. Chem. A*, 2018, **6**, 24441–24451.
- L. Sun, J. Liu, Y. Zhao, J. Xu and Y. Li, *Carbon*, 2019, **145**, 352–358.
- A. M. Ahmed, A. I. Ahmed, M. I. Amin and U. A. El-Banna, *Desalin. Water Treat.*, 2015, **54**(6), 1694–1707.
- M. Gao, L. Zhu, C. K. Peh and G. W. Ho, *Energy Environ. Sci.*, 2019, **12**, 841–864.
- F. Zhao, X. Zhou, Y. Shi, X. Qian, M. Alexander, X. Zhao, S. Mendez, R. Yang, L. Qu and G. Yu, *Nat. Nanotechnol.*, 2018, **13**, 489–495.
- K. Kim, S. Yu, S.-Y. Kang, S.-T. Ryu and J.-H. Jang, *Desalination*, 2019, **469**, 114091.
- Y. Shi, R. Li, Y. Jin, S. Zhuo, L. Shi, J. Chang, S. Hong, K.-C. Ng and P. Wang, *Joule*, 2018, **2**, 1171–1186.



- 22 P. Mu, W. Bai, Z. Zhang, J. He, H. Sun, Z. Zhu, W. Liang and A. Li, *J. Mater. Chem. A*, 2018, **6**, 18183–18190.
- 23 B. Yu, Y. Wang, Y. Zhang and Z. Zhang, *Nano Res.*, 2022, **16**, 5610.
- 24 J. Chen, J. Feng, Z. Li, P. Xu, X. Wang, W. Yin, M. Wang, X. Ge and Y. Yin, *Nano Lett.*, 2019, **19**, 400–407.
- 25 F. He, X. Wu, J. Gao and Z. Wang, *J. Mater. Chem. A*, 2021, **9**, 27121–27139.
- 26 Y. Isaka, Y. Kondo, Y. Kawase, Y. Kuwahara, K. Mori and H. Yamashita, *Chem. Commun.*, 2018, **54**, 9270–9273.
- 27 W. Zhan, L. Sun and X. Han, *Nano-Micro Lett.*, 2019, **11**, 1.
- 28 S. N. Zhao, G. Wang, D. Poelman and P. Van Der Voort, *Molecules*, 2018, **23**, 2947.
- 29 A. Dhakshinamoorthy, Z. Li and H. Garcia, *Chem. Soc. Rev.*, 2018, **47**, 8134–8172.
- 30 J. Zhu, P.-Z. Li, W. Guo, Y. Zhao and R. Zou, *Coord. Chem. Rev.*, 2018, **359**, 80–101.
- 31 K. Yue, X. Zhang, S. Jiang, J. Chen, Y. Yang, F. Bi and Y. Wang, *J. Mol. Liq.*, 2021, **335**, 116108.
- 32 J. D. Xiao and H. L. Jiang, *Acc. Chem. Res.*, 2019, **52**, 356–366.
- 33 H. Yang, M. Zhao, J. Zhang, J. Ma, P. Wu, W. Liu and L. Wen, *J. Mater. Chem. A*, 2019, **7**, 20742–20749.
- 34 M. Tahir, B. Tahir, N. A. S. Amin and A. Muhammad, *Energy Convers. Manage.*, 2016, **119**, 368–378.
- 35 G. Ni, G. Li, S. V. Boriskina, H. Li, W. Yang, T. Zhang and G. Chen, *Nat. Energy*, 2016, **1**, 16126.
- 36 M. Kadhom, W. Hu and B. Deng, *Membranes*, 2017, **7**(2), 31.
- 37 Y. Liu, J. Lou, M. Ni, C. Song, J. Wu, N. P. Dasgupta, P. Tao, W. Shang and T. Deng, *ACS Appl. Mater. Interfaces*, 2016, **8**, 772–779.
- 38 A. Muller, S. Peglow, M. Karnahl, A. Kruth, H. Junge, V. Bruser and C. Scheu, *Nanomaterials*, 2018, **8**(7), 502.
- 39 S. A. El-Hakam, F. T. Alshorifi, R. S. Salama, S. Gamal, W. S. A. El-Yazeed, A. A. Ibrahim and A. I. Ahmed, *J. Mater. Res. Technol.*, 2022, **18**, 1963–1976.
- 40 A. A. Ibrahim, R. S. Salama, S. A. El-Hakam, A. S. Khder and A. I. Ahmed, *Colloids Surf., A*, 2021, **631**, 127753.
- 41 A. A. Ibrahim, S. L. Ali, M. S. Adly, S. A. El-Hakam, S. E. Samra and A. I. Ahmed, *RSC Adv.*, 2021, **11**, 37276–37289.
- 42 A. A. Ibrahim, R. S. Salama, S. A. El-Hakam, A. S. Khder and A. I. Ahmed, *Colloids Surf., A*, 2021, **616**, 126361.
- 43 M. M. Kaid, A. Gebreil, S. A. El-Hakam, A. I. Ahmed and A. A. Ibrahim, *RSC Adv.*, 2020, **10**, 15586–15597.
- 44 W. S. Abo El-Yazeed, S. A. El-Hakam, A. A. Salah and A. A. Ibrahim, *J. Photochem. Photobiol., A*, 2021, **417**, 113362.
- 45 W. S. A. Abo El-Yazeed and A. I. Ahmed, *RSC Adv.*, 2019, **9**(33), 18803–18813.
- 46 X. Han, L. Zang, S. Zhang, T. Dou, L. Li, J. Yang, L. Sun, Y. Zhang and C. Wang, *RSC Adv.*, 2020, **10**, 2507–2512.
- 47 C. Shen, Y. Zhu, X. Xiao, X. Xu, X. Chen and G. Xu, *ACS Appl. Mater. Interfaces*, 2020, **12**, 35142–35151.
- 48 V. Aliamradni, S. S. Abolmaali and S. Borandeh, *J. Nanostruct.*, 2019, **9**, 402–413.
- 49 H. Ma, H. Geng, B. Yao, M. Wu, C. Li, M. Zhang, F. Chi and L. Qu, *ACS Nano*, 2019, **13**, 9161–9170.
- 50 M. Shaban, M. Rabia, W. Fathallah, N. A. El-Mawgoud, A. Mahmoud, H. Hussien and O. Said, *J. Polym. Environ.*, 2017, **26**, 434–442.
- 51 P. Boomi, G. P. Poorani, S. Palanisamy, S. Selvam, G. Ramanathan, S. Ravikumar, H. Barabadi, H. G. Prabu, J. Jeyakanthan and M. Saravanan, *J. Cluster Sci.*, 2019, **30**, 715–726.
- 52 R. Bushra, T. Arfin, M. Oves, W. Raza, F. Mohammad, M. A. Khan, A. Ahmad, A. Azam and M. Muneer, *New J. Chem.*, 2016, **40**, 9448–9459.
- 53 Y. Zou, X. Chen, W. Guo, X. Liu and Y. Li, *ACS Appl. Energy Mater.*, 2020, **3**, 2634–2642.
- 54 X. Han, X. Yang, G. Liu, Z. Li and L. Shao, *Chem. Eng. Res. Des.*, 2019, **143**, 90–99.
- 55 J. O. Olowoyo, U. Saini, M. Kumar, H. Valdés, H. Singh, M. O. Omorogie, J. O. Babalola, A. V. Vorontsov, U. Kumar and P. G. Smirniotis, *J. CO₂ Util.*, 2020, **42**, 101300.
- 56 Z. Li, G. Che, W. Jiang, L. Liu and H. Wang, *RSC Adv.*, 2019, **9**, 33238–33245.
- 57 A. Rahmani, H. B. M. Emrooz, S. Abedi and A. Morsali, *Mater. Sci. Semicond. Process.*, 2018, **80**, 44–51.
- 58 B. Bakbolat, C. Daulbayev, F. Sultanov, R. Beissenov, A. Umirzakov, A. Mereke, A. Bekbaev and I. Chuprakov, *Nanomaterials*, 2020, **10**(9), 1790.
- 59 Y. Zhao, W. Cai, J. Chen, Y. Miao and Y. Bu, *Front. Chem.*, 2019, **7**, 789.
- 60 Q. Wen, L. Ma, C. Wang, B. Wang, R. Han, C. Hao and K. Chen, *RSC Adv.*, 2019, **9**, 14520–14530.
- 61 S. Miao, H. Zhang, S. Cui and J. Yang, *Chemosphere*, 2020, **257**, 127123.
- 62 Z. Yang, X. Xu, X. Liang, C. Lei, Y. Cui, W. Wu, Y. Yang, Z. Zhang and Z. Lei, *Appl. Catal., B*, 2017, **205**, 42–54.
- 63 Z. Yang, J. Ding, J. Feng, C. He, Y. Li, X. Tong, X. Niu and H. Zhang, *Appl. Organomet. Chem.*, 2018, **32**(4), 4285.
- 64 Z. Dargahi, H. Asgharzadeh and H. Maleki-Ghaleh, *Ceram. Int.*, 2018, **44**, 13015–13023.
- 65 B. Singaram, J. Jeyaram, R. Rajendran, P. Arumugam and K. Varadharajan, *Ionics*, 2018, **25**, 773–784.
- 66 Y. P. Putra, S. Wahyuningsih, A. H. Ramelan and R. Hidayat, *Mater. Res. Express*, 2019, **6**(12), 126207.
- 67 S. V. Nipane, S.-W. Lee, G. S. Gokavi and A. N. Kadam, *J. Mater. Sci.: Mater. Electron.*, 2018, **29**, 16553–16564.
- 68 M. Shabani-Nooshabadi and F. Karimian-Taheri, *RSC Adv.*, 2015, **5**, 96601–96610.
- 69 Y. Zhang, J. Liu, Y. Zhang, J. Liu and Y. Duan, *RSC Adv.*, 2017, **7**, 54031–54038.
- 70 P. Fan, Z. Fan, F. Huang, J. Yang, F. Chen, Z. Fei and M. Zhong, *Ind. Eng. Chem. Res.*, 2019, **58**, 3893–3901.
- 71 S. Lyu, Y. Chen, S. Han, L. Guo, Z. Chen, Y. Lu, Y. Chen, N. Yang and S. Wang, *RSC Adv.*, 2018, **8**, 13191–13199.
- 72 M. S. Adly, S. M. El-Dafrawy, A. A. Ibrahim, S. A. El-Hakam and M. S. El-Shall, *RSC Adv.*, 2021, **11**, 13940–13950.
- 73 J. A. Bobb, A. A. Ibrahim and M. S. El-Shall, *ACS Appl. Nano Mater.*, 2018, **1**, 4852–4862.
- 74 P. Ahuja, S. K. Ujjain, I. Arora and M. Samim, *ACS Omega*, 2018, **3**, 7846–7855.



- 75 A. Viswanathan and A. N. Shetty, *Electrochim. Acta*, 2018, **289**, 204–217.
- 76 H. Wang, X. Yuan, Y. Wu, G. Zeng, X. Chen, L. Leng, Z. Wu, L. Jiang and H. Li, *J. Hazard. Mater.*, 2015, **286**, 187–194.
- 77 N. Popovych, P. Kyriienko, S. Soloviev, R. Baran, Y. Millot and S. Dzwigaj, *Phys. Chem. Chem. Phys.*, 2016, **18**, 29458–29465.
- 78 P. Bazant, I. Kuritka, L. Munster and L. Kalina, *Cellulose*, 2015, **22**, 1275–1293.
- 79 F. S. Awad, H. D. Kiriarachchi, K. M. AbouZeid, Ü. Özgür and M. S. El-Shall, *ACS Appl. Energy Mater.*, 2018, **1**, 976–985.
- 80 Y. Zhang, H. Yan, X. Wang, Z. Zhang, F. Liu, S. Tu and X. Chen, *RSC Adv.*, 2022, **12**, 28997–29002.
- 81 G. Chen, J. Sun, Q. Peng, Q. Sun, G. Wang, Y. Cai, X. Gu, Z. Shuai and B. Z. Tang, *Adv. Mater.*, 2020, **32**, e1908537.
- 82 M. Fuseini, M. M. Y. Zaghloul, M. F. Elkady and A. H. El-Shazly, *J. Mater. Sci.*, 2022, **57**, 6085–6101.
- 83 W. Jiang, Z. Li, C. Liu, D. Wang, G. Yan, B. Liu and G. Che, *J. Alloys Compd.*, 2021, **854**, 157166.
- 84 B. Zhai, Y. Chen and Y. Liang, *J. Nanopart. Res.*, 2019, **21**, 1–15.
- 85 Y. Teng, S. Li, C. Xue, H. Zhang, L. Zhu and Y. Tang, *Adv. Polym. Technol.*, 2020, **2020**, 1–9.
- 86 D. A. Kospa, A. I. Ahmed, S. E. Samra and A. A. Ibrahim, *RSC Adv.*, 2021, **11**, 15184–15194.
- 87 Y. A. Younes, D. A. Kospa, R. S. Salama, A. I. Ahmed and A. A. Ibrahim, *Desalination*, 2023, **550**, 116377.
- 88 Y. Fang, Y. Ma, M. Zheng, P. Yang, A. M. Asiri and X. Wang, *Coord. Chem. Rev.*, 2018, **373**, 83–115.
- 89 R. Li, W. Zhang and K. Zhou, *Adv. Mater.*, 2018, **30**, e1705512.
- 90 B. Shao, X. Wu, Y. Wang, T. Gao, Z.-Q. Liu, G. Owens and H. Xu, *J. Mater. Chem. A*, 2020, **8**, 24703–24709.
- 91 H.-S. Kim, H. L. Hobbs, L. Wang, M. J. Rutten and C. C. Wamser, *Synth. Met.*, 2009, **159**, 1313–1318.
- 92 P. Karthik, R. Vinoth, P. Zhang, W. Choi, E. Balaraman and B. Neppolian, *ACS Appl. Energy Mater.*, 2018, **1**, 1913–1923.
- 93 R. Mukherjee, R. Sharma, P. Saini and S. De, *Environ. Sci.: Water Res. Technol.*, 2015, **1**, 893–904.
- 94 J. Yang, J. Choi, D. Bang, E. Kim, E. K. Lim, H. Park, J. S. Suh, K. Lee, K. H. Yoo, E. K. Kim, Y. M. Huh and S. Haam, *Angew. Chem., Int. Ed. Engl.*, 2011, **50**, 441–444.
- 95 Y. Xiao, X. Wang, C. Li, H. Peng, T. Zhang and M. Ye, *J. Environ. Chem. Eng.*, 2021, **9**(1), 105010.
- 96 G. Liu, T. Chen, J. Xu, G. Yao, J. Xie, Y. Cheng, Z. Miao and K. Wang, *Cell Rep. Phys. Sci.*, 2021, **2**, 100310.
- 97 D. A. Kospa, A. Gebreil, S. A. El-Hakam, A. I. Ahmed and A. A. Ibrahim, *J. Mater. Res. Technol.*, 2023, **23**, 2924–2939.
- 98 A. G. Saad, A. Gebreil, D. A. Kospa, S. A. El-Hakam and A. A. Ibrahim, *Desalination*, 2022, **535**, 115824.
- 99 X. Wang, G. Ou, N. Wang and H. Wu, *ACS Appl. Mater. Interfaces*, 2016, **8**, 9194–9199.
- 100 Y. Fu, G. Wang, X. Ming, X. Liu, B. Hou, T. Mei, J. Li, J. Wang and X. Wang, *Carbon*, 2018, **130**, 250–256.
- 101 H. Li, Y. He, Z. Liu, B. Jiang and Y. Huang, *Energy*, 2017, **139**, 210–219.
- 102 J. Huang, Y. He, Y. Hu and X. Wang, *Energy*, 2018, **165**, 1282–1291.
- 103 G. Wang, Y. Fu, A. Guo, T. Mei, J. Wang, J. Li and X. Wang, *Chem. Mater.*, 2017, **29**, 5629–5635.
- 104 A. K. Sadoun, A. Gebreil, R. M. Eltabey, D. A. Kospa, A. I. Ahmed and A. A. Ibrahim, *RSC Adv.*, 2022, **12**, 28843–28852.
- 105 L. Ying, H. Zhu, H. Huang, X. Qu, C. Wang, X. Wang, F. Duan, S. Lu and M. Du, *ACS Appl. Energy Mater.*, 2021, **4**, 3563–3572.
- 106 X. Li, D. Yue, F. Liu, J. Yu, B. Li, D. Sun and X. Ma, *Korean J. Chem. Eng.*, 2023, **40**, 223–234.
- 107 V. Kashyap and H. Ghasemi, *J. Mater. Chem. A*, 2020, **8**, 7035–7065.
- 108 I. Akin, E. Zor, H. Bingol and M. Ersoz, *J. Phys. Chem. B*, 2014, **118**, 5707–5716.
- 109 J. Chen, F. Cheng, D. Luo, J. Huang, J. Ouyang, A. Nezamzadeh-Ejhieh, M. S. Khan, J. Liu and Y. Peng, *Dalton Trans.*, 2022, **51**, 14817–14832.
- 110 A. Mirmohseni, M. Rastgar and A. Olad, *J. Appl. Polym. Sci.*, 2019, **136**(23), 47629.
- 111 S. P. Deshmukh, A. G. Dhodamani, S. M. Patil, S. B. Mullani, K. V. More and S. D. Delekar, *ACS Omega*, 2020, **5**, 219–227.
- 112 K. M. M. Ismayil, A. Varghese and R. Antony, *J. Elastomers Plast.*, 2019, **52**, 103–116.