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# Bio-based sunflower carbon/polyethylene glycol shape-stabilized phase change materials for thermal energy storage†

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The exploitation of shape-stabilized phase change materials with high thermal conductivity and energy storage capacity is an effective strategy for improving energy efficiency. In this work, sunflower stem carbon/polyethylene glycol (SS-PEG) and sunflower receptacle carbon/polyethylene glycol (SR-PEG) shape-stabilized phase change materials, utilizing sunflower stem and receptacle biomass carbon with high specific surface area and pore volume obtained by carbonization as frameworks and polyethylene glycol as an energy storage material, were prepared by the vacuum impregnation method. The ability to load polyethylene glycol into the pore structure of carbon materials in different sunflower parts was mainly investigated, and the micro-morphology, compositional structure and thermal properties were characterized and analyzed using SEM, IR spectroscopy, XRD, DSC and TG techniques. The results showed that the carbonized sunflower stems maintained the sieve pore structure, and the carbonized sunflower receptacle was a macroporous structure containing a large number of three-dimensional interconnections. At the same time, the interaction between polyethylene glycol and each carbon material occurred through physisorption. The melting enthalpies of SS-PEG and SR-PEG shape-stabilized phase change materials were 153.4 J g<sup>-1</sup> and 171.5 J g<sup>-1</sup>, respectively, and the loading rates reached 81.9% and 91.5%, with initial thermal decomposition temperatures ( $T_{5\%}$ ) of 344 °C and 368 °C.

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## 1. Introduction

With the rapid development of the global society and economy, energy consumption and environmental pollution from fossil energy sources such as coal have been increasing.<sup>1–3</sup> To meet human needs and achieve the goal of energy conservation, emission reduction, and improvement of environmental pollution, the future global energy trend is to reduce the use of disposable traditional energy sources such as coal, oil, and natural gas, develop new energy sources and promote the development of renewable energy sources. Thermal energy storage (TES) technology is an effective strategy to improve energy utilization, which mainly consists of four types: sensible heat, latent heat, thermochemical, and sorption heat storage.<sup>4,5</sup> Among the various thermal energy storage methods, latent heat storage (LHS) based on phase change materials (PCMs) is an efficient energy storage technology with high energy storage density within the phase change temperature interval of the

material, which utilizes the change of physical state of phase change materials to absorb or release energy and realizes high-efficiency energy storage.<sup>6,7</sup>

At present, phase change materials with considerable energy storage density have become a research hotspot in the industry as energy-saving materials and have been widely used in the fields of solar energy systems,<sup>8</sup> the automobile industry,<sup>9</sup> astronautics and aeronautics,<sup>10</sup> energy-saving buildings,<sup>11</sup> and intelligent textiles.<sup>12</sup> These are mainly divided into solid–solid phase change materials and solid–liquid phase change materials according to the classification of the phase transition form of PCMs. Solid–solid phase change materials are primarily organic materials with good application prospects in thermal energy storage due to their advantages of no-phase separation, small volume change, anti-leakage, and excellent mechanical properties,<sup>13,14</sup> mainly including polyol phase change materials, layered chalcogenides and polymers, and their latent heat of phase transition is smaller than that of solid–liquid phase change materials while their phase transition temperature is relatively high. Solid–liquid phase change materials can be divided into organic phase change materials, inorganic phase change materials, and eutectic phase change materials according to their chemical compositions, including paraffin, fatty acids, alcohols, crystalline hydrated salts, molten salts, and eutectic hydrated salts, which have been widely used due to

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their sizeable latent heat values, many types and small volume changes.<sup>15–17</sup> Among them, polyethylene glycol (PEG) is an organic phase change material with the advantages of non-toxicity, favorable biocompatibility, and excellent thermal stability.<sup>14</sup> However, the fluidity and the susceptibility to leakage of the liquid phase of polyethylene glycol during the phase change process and low thermal conductivity limit their applications.<sup>1,18,19</sup> Confining polyethylene glycol to porous-based supporting materials such as polymers,<sup>20,21</sup> foam metal,<sup>22,23</sup> and porous carbon materials<sup>19,24</sup> to prepare shape-stabilized phase change materials is an effective strategy to address the above drawbacks.

Biomass porous carbon materials have attracted widespread attention and are widely used in many fields, such as energy storage,<sup>25,26</sup> supercapacitors,<sup>27,28</sup> wastewater treatment,<sup>29,30</sup> and electromagnetic shielding,<sup>31,32</sup> because of their vast source, low cost, and renewability, and are synthesized from biomass through thermochemical degradation processes such as gasification, hydrothermal carbonization, and high-temperature pyrolysis under anoxic or micro-oxygen conditions.<sup>33</sup> Biomass is a plant resource composed of organic carbon,<sup>34</sup> which primarily includes agricultural by-products such as straw, rice husk, corn kernel, fruit shells, and bamboo wood, and its chemical composition is mainly cellulose, hemicellulose and lignin, some of which contain a small amount of silica.<sup>35</sup> Compared with ordinary carbon materials, biomass porous carbon materials have apparent advantages: cross-linked three-dimensional porous skeleton structure, high specific surface area, strong corrosion and heat resistance, and high thermal and electrical conductivity. Currently, biomass porous carbon materials are prepared using carbonization, activation and template methods. Among them, the direct carbonization method has the advantages of simple operation and low cost, which involves thermal decomposition at a high temperature in an inert gas ( $N_2$  or He) atmosphere, and the non-carbon elements and volatile components are removed through various physical and chemical reactions, resulting in the formation of porous carbon.<sup>36</sup> Wei *et al.*<sup>37</sup> prepared succulent-based carbon aerogel (SCA) by carbonizing the leaves of succulent plants and introducing molten paraffin into the SCA to obtain a shape-stabilized phase change material. The results showed that the maximum loading of SCA to paraffin was up to 95%, and the melting enthalpy of the composite phase change material was  $130.1 \text{ J g}^{-1}$ , which was only 1.8% lower than that of pure paraffin. Ji *et al.*<sup>38</sup> prepared a 3D porous network carbon aerogel with “soft-rigid” network structures by direct carbonization of pomelo peels as a support matrix loading paraffin wax to acquire a composite phase change material, which showed a significant latent heat storage capacity of  $159.9 \text{ J g}^{-1}$  and exhibited excellent thermal reliability after 25 frequent heating/cooling cycles. The composite phase change materials exhibit ultra-low density, excellent thermal stability, enhanced thermal and electrical conductivity, shape stability, and leakage resistance.

Sunflower stems and receptacles have a good pore structure, which provides a circulation path for the carbonized skeleton loaded with phase change materials. Here, we utilize

carbonized sunflower stems and receptacles loaded with polyethylene glycol for the first time with a view to preparing two shape-stabilized phase change materials. In this paper, sunflower stem biomass carbon (SSC) and sunflower receptacle biomass carbon (SRC) with high specific surface area and pore volume were prepared from sunflower biomass by direct carbonization, and sunflower stem carbon/polyethylene glycol and sunflower containing carbon/polyethylene glycol shape-stabilized phase change materials were prepared from polyethylene glycol loaded into the two biomass porous carbon materials using vacuum impregnation method. The loading capacity of the porous structure of the two biomass porous carbon materials for polyethylene glycol was investigated, and their thermal storage properties were also explored. Thus, this work opens up new avenues for efficient energy utilization.

## 2. Experimental methods

### 2.1. Materials

Sunflowers were collected from the Sichuan Province of China. Polyethylene glycol (PEG,  $M_n = 4000$ ) was provided by Bayer Co., Ltd (China).

### 2.2. Preparation of sunflower biomass carbon

The purchased raw materials were washed with deionized water and placed in a sunny environment at room temperature for 24 h. The excess water was wiped off using paper towels, and each part of the sunflower (stem, receptacle, leaves, and petal) was separated appropriately before being placed in a freeze-drying oven at  $-70^\circ\text{C}$  for 4 h and then freeze-dried for 48 h. The lyophilized and dehydrated sunflower parts were heated up to  $1000^\circ\text{C}$  under the atmosphere of  $N_2$  in a tube furnace at  $5^\circ\text{C min}^{-1}$  and pyrolyzed for 3 h at this temperature to obtain SSC, SRC, sunflower leaf biomass carbon (SLC) and sunflower petal biomass carbon (SPC), as shown in Fig. 1.

### 2.3. Preparation of shape-stabilized phase change materials

The PEG was heated to  $80^\circ\text{C}$  in an oven to melt into the liquid state. Then, two samples, SSC and SRC, were immersed into the molten PEG under a vacuum of  $-0.1 \text{ MPa}$  for 4 h to allow the PEG to be adsorbed by the activated charcoal and expel the internal bubbles. Moreover, the samples were removed and placed on filter paper at  $80^\circ\text{C}$ , the excess PEG adsorbed on the surface of the activated charcoal was rolled off until there was no leakage from the filter paper, and the shape-stabilized phase change materials were obtained. The flowchart for preparing shape-stabilized phase change materials is shown in Fig. 1.

### 2.4. Characterizations

The micro-morphology of samples was investigated by field emission scanning electron microscopy (SEM, JSM-5900LV, Japan). The porous structure of SSC and SRC samples was analyzed using Brunauer–Emmett–Teller (BET, Micromeritics ASAP2460, American). The graphitization degrees of SSC and SRC samples were characterized by Raman spectra (inVia Reflex, American). The chemical structures of samples were



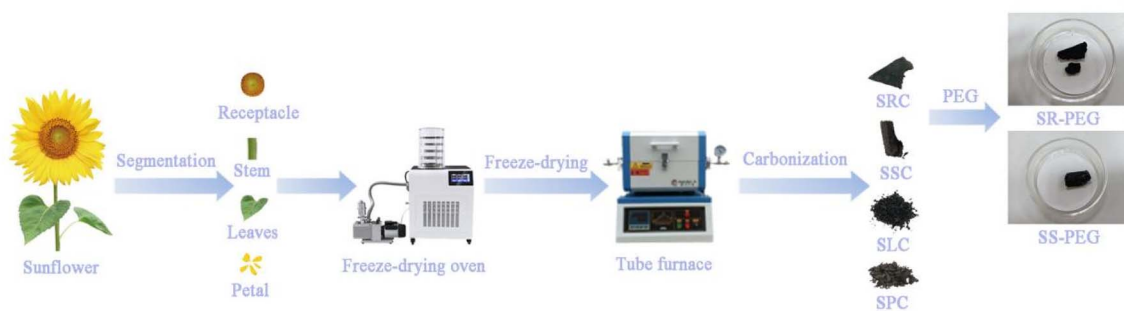


Fig. 1 The flowchart for the preparation of shape-stabilized phase change materials.

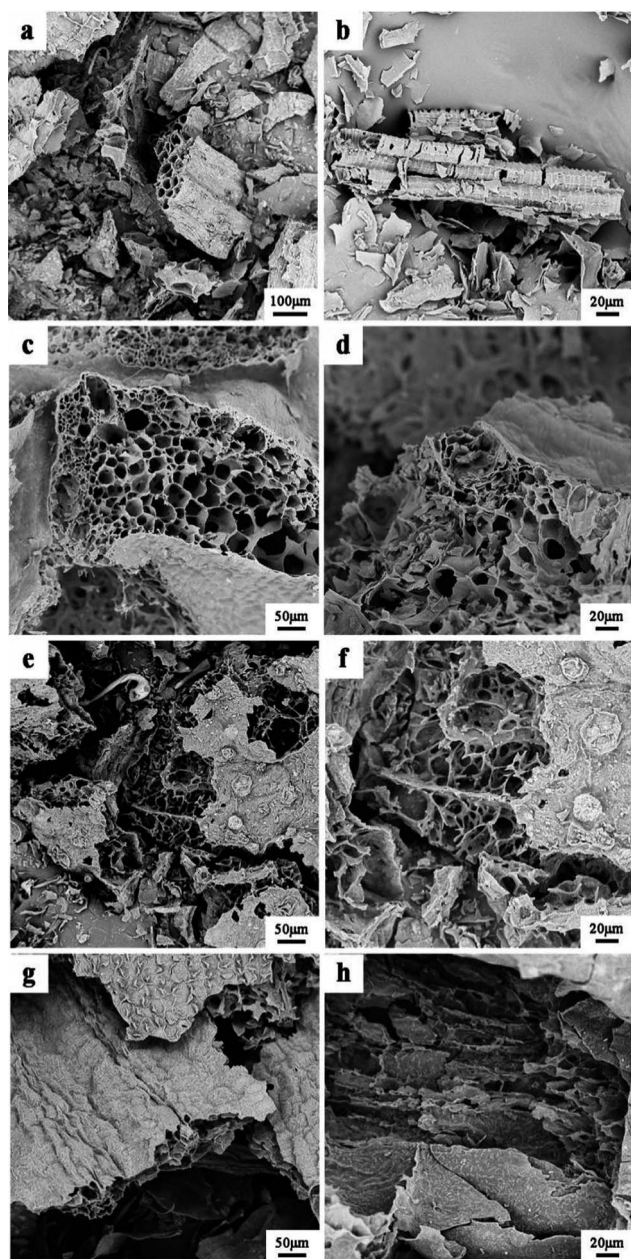


Fig. 2 SEM images of primary carbon in various parts of sunflower: (a and b) SSC; (c and d) SRC; (e and f) SLC; (g and h) SPC.

studied by Fourier transform infrared spectroscopy (FT-IR, Nicolet 50, American) in the infrared wavenumber from 4000 to 400  $\text{cm}^{-1}$ , and X-ray diffraction (XRD, Ultima IV, Japan, Cu Ka radiation at 40 kV and 50 mA). The thermal properties of samples were analyzed by differential scanning calorimetry (DSC, TA Q250, American). A 3–5 mg sample was taken and sealed in an aluminum crucible and placed in the apparatus, and the sample was heated from 10  $^{\circ}\text{C}$  to 90  $^{\circ}\text{C}$  at a heating rate of 10  $^{\circ}\text{C min}^{-1}$  and then cooled down from 90  $^{\circ}\text{C}$  to 10  $^{\circ}\text{C}$  at a cooling rate of 10  $^{\circ}\text{C min}^{-1}$  in a nitrogen atmosphere. In order to ensure the reliability of data, three specimens were tested for each sample to obtain an average value. The thermal decomposition behavior of samples was observed by thermogravimetric analysis (TGA, TGA2, Sweden) from 35 to 800  $^{\circ}\text{C}$  at a heating rate of 10  $^{\circ}\text{C min}^{-1}$  under  $\text{N}_2$  atmosphere. Thermal conductivity of two shape-stabilized PCMs was conducted by the thermal constant analyzer (Model 2500-OT, Hot disk, Sweden) using a Kapton probe with a radius of 2.001 mm. Solar thermal energy conversion capability was evaluated *via* an infrared thermal imager (FTLR T420, American). The shape stability of the samples was examined by photographing the leakage of the samples before and after 30 min at 80  $^{\circ}\text{C}$ .

### 3. Results and discussion

#### 3.1. Structures of SSC and SRC

Fig. 2 shows the SEM images of the raw carbon of each part of the sunflower. Fig. 2a and b shows the morphology of the pore structure of the sunflower stem biomass carbon material. It could be clearly seen that there was a sieve pore structure inside the SSC, which maintained the pore structure of the raw material, while the pith of the stem in the central layer was more easily broken by pressure. The edge of the lignified structure was relatively more potent and retained some of the original sieve pore structure after extrusion and crushing. Fig. 2c and d presented the morphology of the pore structure of the carbon material of the sunflower receptacle, from which it could be obviously observed that the SRC also retained the morphology of the original material, containing a large number of three-dimensional interconnected macroporous structures, with the pore size of 10–50  $\mu\text{m}$ . Fig. 2e–h show the morphology of the pore structures of the sunflower leaf biomass carbon and the





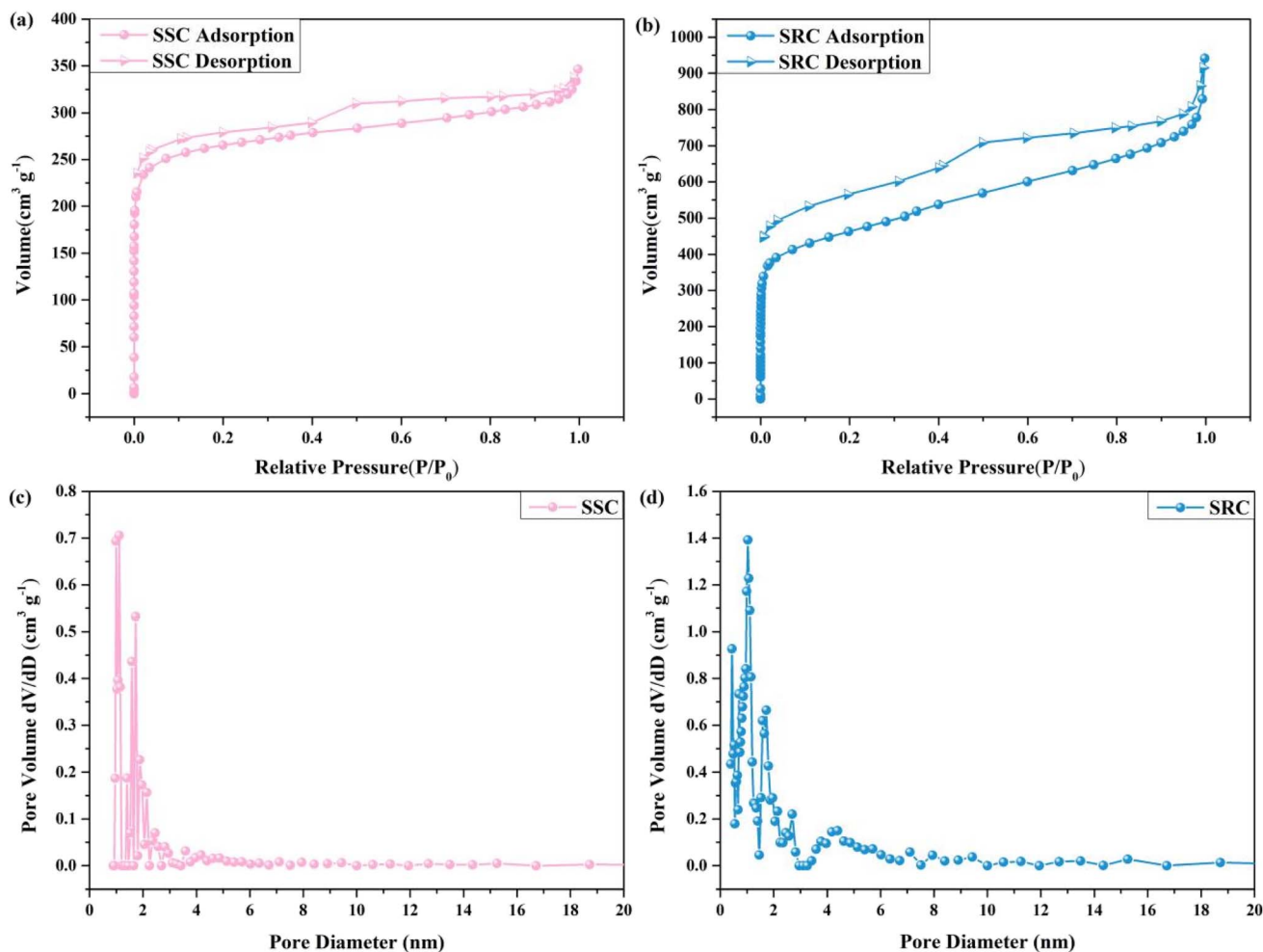


Fig. 3 Nitrogen adsorption/desorption isotherms: (a) SSC and (b) SRC, and DFT desorption pore size distribution: (c) SSC and (d) SRC.

sunflower petal biomass carbon materials, respectively. The pore structure of the SLC material is mainly a three-dimensional interpenetrating irregular pore structure, and the pore structure of the SPC material is a unidirectional pore channel structure. Due to the defects of low yield and fragility of the SLC and SPC materials, although they also have excellent pore structures, only SSC and SRC materials were used as the skeleton support materials when impregnating PEG in the later stage.

In order to assess the pore structure and adsorption capability of SSC and SRC, the specific surface area and pore size of SSC and SRC were analyzed utilizing  $N_2$  adsorption-desorption isotherms and DFT desorption pore size distribution, as shown

in Fig. 3 and Table 1. The isotherms of SSC and SRC were both type IV accompanied by an H4 hysteresis loop according to the IUPAC classification.<sup>39</sup> Fig. 3a and b showed that the adsorption of SSC and SRC increased rapidly in the relatively low-pressure region, indicating the existence of a strong interaction force between SSC and nitrogen as well as between SRC and nitrogen, along with the existence of a microporous structure. In the higher pressure region ( $0.5 < P/P_0 < 0.99$ ), the adsorption of SSC and SRC again increased steeply due to capillary condensation, demonstrating the presence of meso/macropores. Combining Fig. 3 with Table 1, the specific surface area of SSC and SRC was  $1016.60 \text{ m}^2 \text{ g}^{-1}$  and  $1637.00 \text{ m}^2 \text{ g}^{-1}$ , respectively, and the pore volume of SSC and SRC was  $0.51 \text{ cm}^3 \text{ g}^{-1}$  and  $1.27 \text{ cm}^3 \text{ g}^{-1}$ ,

Table 1 BET surface area and pore volume of SSC and SRC

Samples	BET surface area ( $\text{m}^2 \text{ g}^{-1}$ )	Total pore volume ( $\text{cm}^3 \text{ g}^{-1}$ )	Micro pore volume ( $\text{cm}^3 \text{ g}^{-1}$ )	Mean pore size (nm)
SSC	1016.60	0.51	233.56	2.01
SRC	1637.00	1.27	376.10	3.10



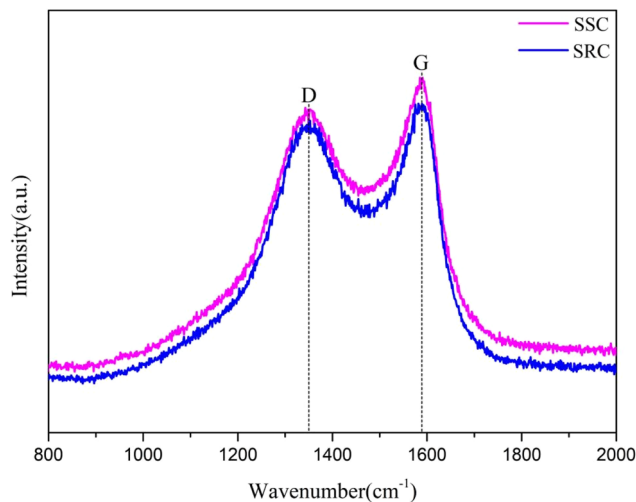


Fig. 4 Raman spectra of SSC and SRC.

which was mainly attributed to the fact that water vapor and carbon dioxide generated from the raw materials can promote the formation of porous structures at high temperatures. Moreover, the average pore size of SSC and SRC was 2.01 nm and 3.10 nm, respectively, showing that SSC and SRC had a multilevel pore structure and their main structure was mesoporous, which contributed to the penetration and loading of organic PCM into SSC and SRC.

The physical properties of the prepared SSC and SRC were investigated by Raman spectroscopy, and the Raman spectra are shown in Fig. 4. The broad peaks at  $1348\text{ cm}^{-1}$  and  $1590\text{ cm}^{-1}$  corresponded to the D and G bands of carbon, respectively, where the relative intensity of the D band represented the degree of defects in the graphite properties, which was known as disordered graphite.<sup>40</sup> The G band correlated with the phonon modes of graphite and the formation of graphitic carbon and was used to characterize the  $\text{sp}^2$  bond structure of the carbon material. The graphite properties of the samples were evaluated using the ratio of the intensity of the G band to the peak intensity of the D band ( $I_{\text{G}}/I_{\text{D}}$ ), and the  $I_{\text{G}}/I_{\text{D}}$  values calculated here are 1.10 and 1.06, respectively, indicating that SSC and SRC had high graphitic structures. Meanwhile, the presence of many structural defects in the two bio-carbon materials implied that some of the carbon in the samples was in an amorphous state due to carbonization, and this amorphousness contributed to the enhancement of the pore properties of the carbon materials, which further improved the adsorption capacity of the organic phase-change materials, and thus enhanced their suitability as a support framework for thermal energy storage materials.<sup>41</sup>

### 3.2. Structures of shape-stabilized PCMs

Two shape-stabilized phase change materials, SS-PEG and SR-PEG, were obtained by loading PEG into SSC and SRC support skeletons through vacuum impregnation, and the morphology and structure of these two materials are shown in Fig. 5. In Fig. 5a, it was observed that the polyethylene glycol had been

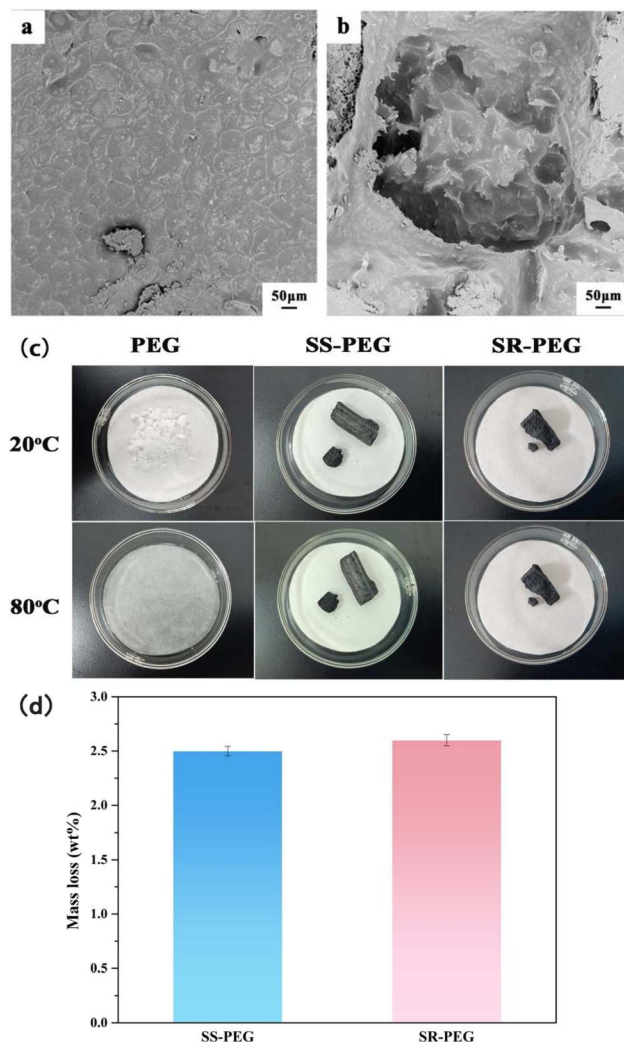


Fig. 5 SEM images of shape-stabilized phase change materials: (a) SS-PEG and (b) SR-PEG; (c) leakage macroscopic morphology of PEG, SS-PEG and SR-PEG under  $80\text{ }^{\circ}\text{C}$  for 30 min; (d) the leakage rate of SS-PEG and SR-PEG after 100 thermal cycles.

completely filled into the sunflower stem carbon material, which suggested that the sieve pore structure of the sunflower stem carbon material facilitated the PEG infiltration and prevented the leakage of PEG through the physisorption force of the pore structure of this material. Fig. 5b found that when the sunflower receptacle carbon material adsorbed polyethylene glycol, the large pores acted as transportation conduits and mesopores and micropores on the pore walls acted as physical adsorption sites to limit the flow of PEG in the liquid state at elevated temperatures. In order to test the shape stabilizing ability of the two stereotyped phase change materials of SS-PEG and SR-PEG, PEG, SS-PEG and SR-PEG at a temperature of  $20\text{ }^{\circ}\text{C}$  were used as the comparison samples and heated to  $80\text{ }^{\circ}\text{C}$  for 30 min using a temperature higher than the melting point of PEG, which was verified by macroscopically observing the dry and wet state of the filter paper, as shown in Fig. 5c. When the temperature was  $20\text{ }^{\circ}\text{C}$ , PEG, SS-PEG and SR-PEG were all in a solid state, and after being heated to  $80\text{ }^{\circ}\text{C}$  for 30 min, PEG was transformed from the

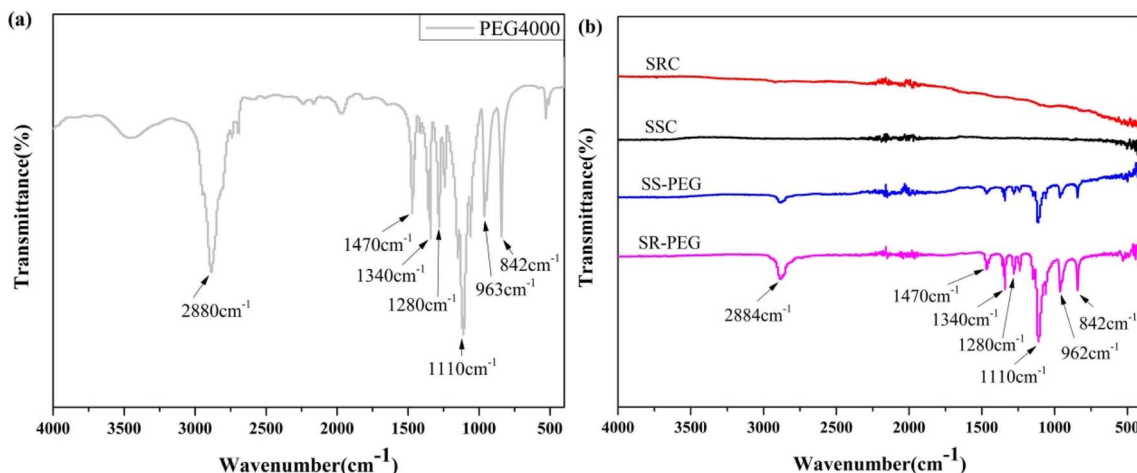


Fig. 6 Infrared spectra of PEG4000, various carbon materials and shape-stabilized phase change materials: (a) PEG4000; (b) SSC, SRC, SS-PEG and SR-PEG.

solid state to the liquid state, SS-PEG and SR-PEG were still in solid state and the filter paper did not show any wetting phenomenon, which implied that SS-PEG and SR-PEG were macroscopically two kinds of shape-stabilized phase change materials. To further study the shape stabilization of the materials, the leakage of the composites was determined by heating the samples cyclically 100 times, and the results are shown in Fig. 5d. After 100 heating cycles, the mass loss of SS-PEG and SR-PEG was 2.49 wt% and 2.64 wt%, respectively, which indicated that the two composites possessed superior leak-proof properties. This excellent leak-proof property can be attributed to the fact that the multistage pore structures of SSC and SRC have sufficient capillary forces to adsorb PCM. During this period, the shape and size of the composites were observed macroscopically without deformation.

In order to analyze the component structure of the SS-PEG and SR-PEG shape-stabilized phase change materials, the

composition of the samples was investigated utilizing Fourier transform infrared absorption spectroscopy. The infrared spectra of PEG, each carbon material, and the SS-PEG and SR-PEG shape-stabilized phase change materials are shown in Fig. 6. In Fig. 6a, the broad absorption peak of PEG at  $3460\text{ cm}^{-1}$  is attributed to the O–H stretching vibration. The characteristic absorption peak at  $2880\text{ cm}^{-1}$  corresponds to the C–H stretching vibration of methylene, while the absorption peaks at  $1470\text{ cm}^{-1}$  and  $1340\text{ cm}^{-1}$  represent the C–H bending vibration of methylene. The absorption peaks at  $1280\text{ cm}^{-1}$  and  $1110\text{ cm}^{-1}$  refer to the C–O–C stretching vibration, whereas the absorption peaks at  $963\text{ cm}^{-1}$  and  $842\text{ cm}^{-1}$  correspond to the C–O–C bending vibration. As can be observed in Fig. 6b, comparing the infrared spectra of PEG and each carbon material, no new characteristic peaks appeared in the infrared spectra of each of the shape-stabilized phase change materials, which implied that there was no chemical reaction between SSC and PEG as well as SRC and PEG during the preparation of the SS-PEG and SR-PEG shape-stabilized phase change materials. Therefore, it primarily relied on the physical adsorption effect to confine the PEG to the pore structure of each carbon material.<sup>42</sup>

The crystallization behaviors of SS-PEG and SR-PEG were analyzed by XRD, as shown in Fig. 7. The high-intensity diffraction peaks of PEG, SS-PEG, and SR-PEG at  $2\theta = 19.08^\circ$  and  $23.18^\circ$  correspond to the characteristic crystal planes (120) and (112), respectively.<sup>14,43</sup> Compared with PEG, the SS-PEG and SR-PEG shape-stabilized phase change materials showed a weakening trend in each peak, which implied that SSC and SRC had a limiting effect on the crystallization of polyethylene glycol. Meanwhile, no new diffraction peaks were generated in the SS-PEG and SR-PEG shape-stabilized phase change materials, which indicated that the crystal structures of PEG in SSC and SRC were not altered as well as PEG was adsorbed into the porous structures of SSC and SRC through physical interactions, in agreement with the analytical results in Fig. 6.

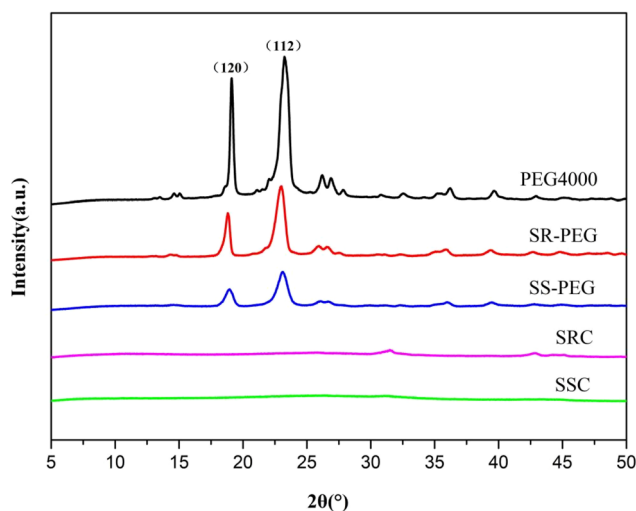


Fig. 7 XRD patterns of PEG4000, SSC, SRC, SS-PEG and SR-PEG.



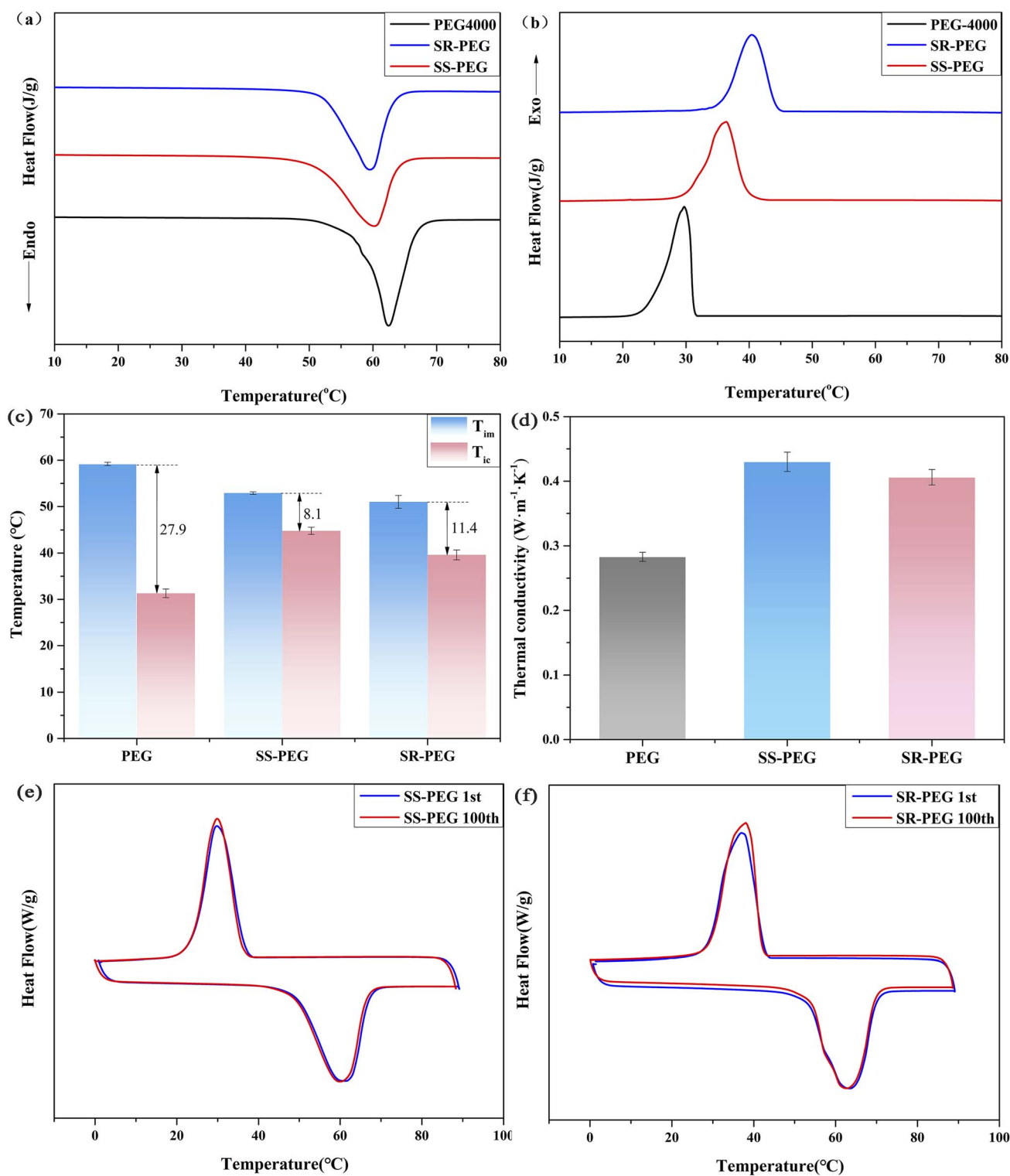


Fig. 8 DSC curves of PEG4000, SS-PEG and SR-PEG: (a) melting curve, (b) crystallization curve; (c) melting and crystallization temperature of PEG4000, SS-PEG and SR-PEG; (d) the thermal conductivity of PEG, SS-PEG, and SR-PEG; DSC curves of composite PCMs after 100 thermal cycles: (e) SS-PEG, (f) SR-PEG.



**Table 2** Enthalpy and phase transition temperature of PEG4000 and various shape-stabilized phase change materials

Samples	$T_{im}$ (°C)	$T_m$ (°C)	$\Delta H_m$ (J g <sup>-1</sup> )	$T_{ic}$ (°C)	$T_c$ (°C)	$\Delta H_c$ (J g <sup>-1</sup> )	$R$ (%)
PEG4000	59.2 ± 0.4	62.5 ± 0.2	187.4 ± 0.3	31.3 ± 1.1	28.5 ± 1.2	174.3 ± 0.5	—
SS-PEG	52.9 ± 0.3	59.7 ± 0.2	153.4 ± 0.7	44.8 ± 0.9	41.4 ± 1.1	150.4 ± 0.4	81.9
SR-PEG	51 ± 1.5	60 ± 1.1	171.5 ± 0.2	39.6 ± 1.2	36.7 ± 0.6	166.5 ± 0.2	91.5

**Table 3** Thermal transition capacity and the loading of recently reported composite PCMs

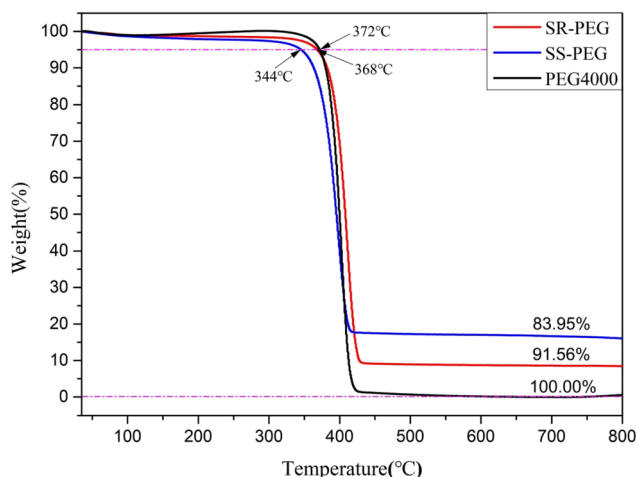
Supporting materials	PCMs	Latent heat (J g <sup>-1</sup> )	Loading (%)	Ref.
Potato derived carbon	PEG4000	91.8	50.0	51
Potato derived carbon	PEG4000	158.8	85.4	52
Eggplant-derived porous carbon	PEG2000	149.0	90.1	53
Hemp-stem-derived biochar	PEG6000	170.44	88.62	54
Corn-cob-derived biochar	PEG6000	121.94	85.19	55
Boron nitride/polypyrrole/wood-based carbon	PEG6000	160	78.1	56
SSC	PEG4000	153.4	81.9	This work
SRC	PEG4000	171.5	91.5	This work

### 3.3. Thermal properties and thermal stability of shape-stabilized PCMs

The thermal performance of SS-PEG and SR-PEG shape-stabilized phase change materials were investigated by DSC, and the results are shown in Fig. 8 and Table 2. The enthalpies of melting and crystallization of PEG were 187.4 J g<sup>-1</sup> and 174.3 J g<sup>-1</sup>, respectively, and the melting and crystallization temperatures were 62.5 °C and 28.5 °C. Compared with PEG, the enthalpies decreased for both SS-PEG and SR-PEG shape-stabilized phase change materials. On the one hand, the addition of SSC and SRC reduced the mass ratio of PEG in the composites, which led to a proportional decrease in the enthalpy of melting/crystallization. On the other hand, the physical adsorption effect of SSC and SRC bound the movement of PEG chain segments and decreased the free movement ability of chain segments. Meanwhile, SSC and SRC also destroyed the

regularity of PEG molecular chains to decrease the crystallinity and melting/crystallization enthalpy of PEG. Compared to pure PEG, the composites showed a different phase transition process, with the melting point decreasing and the crystallization temperature increasing during the heating/cooling cycle. The strong adsorption of SSC and SRC to PEG led to a decrease in the free movement of the PEG molecular chains. Consequently, the energy required for the free movement of PEG molecular chains increased, and the crystallization temperature heightened.

PEG showed very large hysteresis (initial crystallization temperature much below initial melting temperature), which significantly limited its use as a PCM. In Table 2 and Fig. 8c, it was evident that the thermal hysteresis of the composites was improved with the addition of SSC and SRC. According to some studies,<sup>44–46</sup> it is known that the difference in thermal conductivity affected the difference in the hysteresis degree, and under the same conditions, PCM with higher thermal conductivity had a lower melting temperature and higher crystallization temperature, which led to the difference in hysteresis degree. In order to explore the effect of the incorporation of SSC and SRC on the thermal conductivity of the composites, the thermal conductivity of two shape-stabilized PCMs was conducted by the thermal constant analyzer. The thermal conductivity of pure PEG was 0.283 W (m<sup>-1</sup> K<sup>-1</sup>), and the thermal conductivities of SS-PEG and SR-PEG were 0.43 W (m<sup>-1</sup> K<sup>-1</sup>) and 0.406 W (m<sup>-1</sup> K<sup>-1</sup>), respectively. Thermal conductivity of SS-PEG and SR-PEG was significantly enhanced compared to pure polyethylene glycol, which was attributed to the inherently high thermal conductivity of SSC and SRC with higher graphite content, and the three-dimensional interpenetrating porous network structure of SSC and SRC provided a channel for rapid heat transfer. The result is consistent with the literature<sup>47,48</sup> that the addition of biomass porous carbon enhances the thermal conductivity of the matrix material. Therefore, the incorporation of SSC and

**Fig. 9** TGA curves of PEG4000, SS-PEG and SR-PEG.



SRC into the composites increased their thermal conductivity and led to a lower hysteresis degree. The thermal hysteresis of SS-PEG was calculated to be lower than that of SR-PEG. Compared with SR-PEG, the percentage of biochar content of SS-PEG was higher, and the thermal hysteresis was lower with relatively less PEG content per unit mass of the composites, which meant that the thermal hysteresis of the composites decreased with the increase in biochar content.

For the sake of further evaluating the energy storage capacity of SS-PEG and SR-PEG shape-stabilized phase change materials, eqn (1) was used to calculate the loading ratio ( $R$ )<sup>49,50</sup> of sunflower stem and sunflower receptacle biomass carbon (BC) on polyethylene glycol:

$$R = \frac{\Delta H_{m,PCMs/BC}}{\Delta H_{m,PCMs}} \times 100\% \quad (1)$$

$\Delta H_{m,PCMs}$  is the melting enthalpy of pure PEG, and  $\Delta H_{m,PCMs/BC}$  is the melting enthalpy of the prepared SS-PEG and SR-PEG

shape-stabilized phase change materials. From the calculation, it can be seen that the loading ratios of SSC and SRC to PEG are as high as 81.9% and 91.5%, respectively, indicating that SS-PEG and SR-PEG have excellent heat storage capacity compared with other composite phase change materials in Table 3.

Analysis of 100 melting/freezing cycles was performed using DSC to estimate the cyclic stability of SS-PEG and SR-PEG shape-stabilized phase change materials. The 100-cycle test result, shown in Fig. 8e and f, demonstrated that the DSC curve of SS-PEG and SR-PEG after 100 melting/freezing cycles was almost consistent with the first cycle. Compared with the first cycle, the latent heat of SS-PEG and SR-PEG after 50 melting/freezing cycles lost only  $0.59 \text{ J g}^{-1}$  and  $0.84 \text{ J g}^{-1}$ , respectively, which indicated that both SS-PEG and SR-PEG had excellent thermal cycle stability in thermal energy storage and release.

In the practical application process, the thermal stability of composite phase change materials is a crucial parameter. Fig. 9

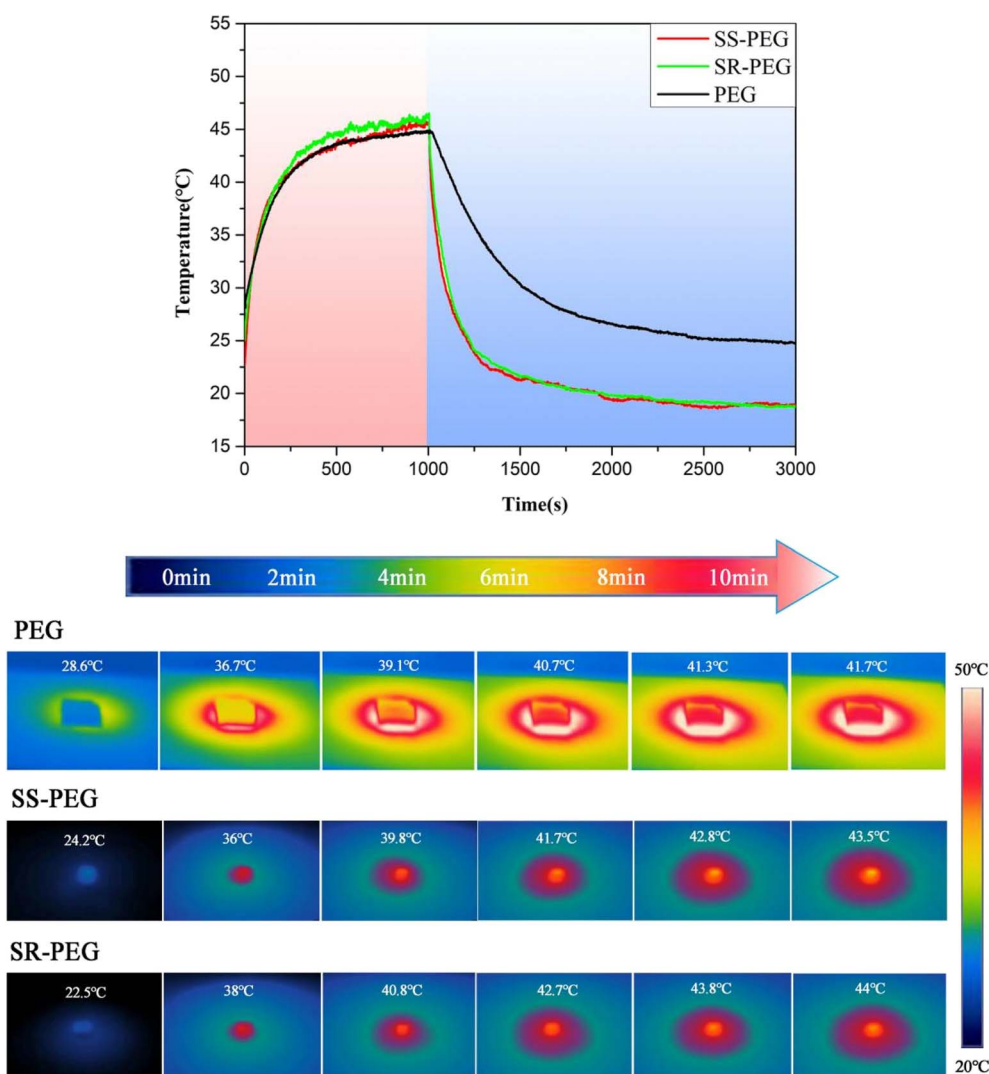


Fig. 10 (a) Temperature curves of PEG, SS-PEG and SR-PEG with time under xenon lamp; (b) infrared thermal images of PEG, SS-PEG and SR-PEG during heating.

shows the thermogravimetric curves of pure PEG, SS-PEG and SR-PEG shape-stabilized phase change materials. As seen in Fig. 9, polyethylene glycol had only one thermogravimetric behavior, and its initial thermal decomposition temperature was 372 °C when the mass loss was 5%, indicating that the polyethylene glycol had good thermal stability. In addition, polyethylene glycol was completely pyrolyzed, and the thermal weight loss rate reached 100% when the temperature increased to 800 °C. Both SS-PEG and SR-PEG also have only one thermal decomposition stage, with initial thermogravimetric loss temperatures ( $T_{5\%}$ ) of 344 °C and 368 °C, respectively. When the temperature reached 418 °C and 436 °C, respectively, the mass loss of SS-PEG and SR-PEG reached the maximum due to the cleavage of PEG molecular chains during the thermal decomposition. Since the weight loss in the whole thermal decomposition process mainly comes from PEG, the load capacity of SSC and SRC to PEG can be further proved by the thermal weight loss rate. When the temperature reached 800 °C, the thermal weight loss rates of SS-PEG and SR-PEG samples were 83.95% and 91.56%, respectively. Combining Table 2 and Fig. 9, the thermal weight loss rate and loading rate of SS-PEG and SR-PEG are roughly the same.

In order to investigate the photothermal conversion ability of SS-PEG and SR-PEG shape-stabilized phase change materials, a xenon lamp was used as a light source to irradiate the samples and an infrared thermal camera was used to capture the temperature changes of the samples. Fig. 10a shows the temperature variation curves of PEG, SS-PEG and SR-PEG with time under a xenon lamp, and Fig. 10b shows the infrared thermograms of PEG, SS-PEG and SR-PEG during the warming process. In Fig. 10b, it can be observed that the temperatures of PEG, SS-PEG and SR-PEG increased by 13.1 °C, 19.3 °C and 21.5 °C, respectively, after irradiation under the same xenon lamp irradiation environment for 10 min. The results demonstrated that SS-PEG and SR-PEG had a faster rate of heating compared to PEG, and the temperatures of SS-PEG and SR-PEG reached higher than those of PEG within the same heating time during the heating process in Fig. 10a and b. Therefore, the addition of SSC and SRC improved the heat transfer ability of SS-PEG and SR-PEG, but SR-PEG had better photothermal conversion compared to SS-PEG.

## 4. Conclusions

Sunflower stem biomass carbon and sunflower receptacle biomass carbon with high specific surface areas and multilayered hierarchical pore structures were prepared by carbonization. The specific surface area of SSC and SRC was 1016.60 m<sup>2</sup> g<sup>-1</sup> and 1637.00 m<sup>2</sup> g<sup>-1</sup>, respectively, and the average pore size was 2.01 nm and 3.10 nm. Sunflower stem/polyethylene glycol and sunflower receptacle/polyethylene glycol shape-stabilized phase change materials were prepared by vacuum impregnation using polyethylene glycol as a phase change material and biomass carbon materials of sunflower stem and sunflower receptacle as the support skeleton. Due to the excellent pore structure of SSC and SRC, the loading rate of SSC and SRC on polyethylene glycol was as high as 81.9% and 91.5%, and the

melting enthalpy of SS-PEG and SR-PEG shape-stabilized phase change materials was 153.4 J g<sup>-1</sup> and 171.5 J g<sup>-1</sup>, respectively. In addition, SS-PEG and SR-PEG had excellent thermal stability and outstanding thermal reliability, and the latent heat remained almost unchanged after 100 cycles. Enhanced heat transfer was observed after the addition of SSC and SRC to PEG. In summary, SS-PEG and SR-PEG shape-stabilized phase change materials have great potential for application in thermal energy storage.

## Data availability

The data supporting this article have been included as part of the ESI.†

## Author contributions

Conceptualization, N. G. and Y. L.; methodology, N. G., J. D., and W. Y.; validation, Y. L. and T. X.; formal analysis, J. D., Y. L., T. X., and C. Y.; investigation, W. Y., B. S., J. L., and B. L.; resources, Y. L. and X. L.; data curation, N. G.; writing – original draft preparation, N. G.; writing – review and editing, Y. L.; supervision, Y. L. and T. X.; project administration, Y. L. All authors have read and agreed to the published version of the manuscript.

## Conflicts of interest

There are no conflicts to declare.

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## References

- 1 Y. Li, B. Wang, W. Zhang, J. Zhao, X. Fang, J. Sun, R. Xia, H. Guo and Y. Liu, *Energy*, 2022, **254**, 124206.
- 2 C. Shao, Y. Zhao and L. Qu, *SusMat*, 2022, **2**, 142–160.
- 3 M. Q. Wu, S. Wu, Y. F. Cai, R. Z. Wang and T. X. Li, *Energy Storage Mater.*, 2021, **42**, 380–417.
- 4 M. Wu, T. Li, Q. He, R. Du and R. Wang, *Energy*, 2022, **239**, 121938.
- 5 X. Chen, P. Cheng, Z. Tang, X. Xu, H. Gao and G. Wang, *Adv. Sci.*, 2021, **8**, 2001274.



- 6 W. Aftab, X. Huang, W. Wu, Z. Liang, A. Mahmood and R. Zou, *Energy Environ. Sci.*, 2018, **11**, 1392–1424.
- 7 G. T. Nguyen, *RSC Adv.*, 2023, **13**, 7621–7631.
- 8 S. Khanna, K. S. Reddy and T. K. Mallick, *Sol. Energy*, 2018, **163**, 591–599.
- 9 V. Talele, P. Thorat, Y. P. Gokhale and V. K. Mathew, *J. Energy Storage*, 2021, **44**, 103482.
- 10 E. Oró, A. D. Gracia, A. Castell, M. M. Farid and L. F. Cabeza, *Appl. Energy*, 2019, **99**, 513e533.
- 11 M. Wu, C. Liu and Z. Rao, *Constr. Build. Mater.*, 2022, **335**, 127400.
- 12 Y. Zheng, H. Oguzlu, A. Baldelli, Y. Zhu, M. Song, A. Pratap-Singh and F. Jiang, *Carbohydr. Polym.*, 2022, **291**, 119583.
- 13 C. Chen, W. Liu, Z. Wang, K. Peng, W. Pan and Q. Xie, *Sol. Energy Mater. Sol. Cells*, 2015, **134**, 80–88.
- 14 N. Gao, T. Tang, H. Xiang, W. Zhang, Y. Li, C. Yang, T. Xia and X. Liu, *Sol. Energy Mater. Sol. Cells*, 2022, **244**, 111831.
- 15 B. Xu, P. Li and C. Chan, *Appl. Energy*, 2015, **160**, 286–307.
- 16 P. Sivasamy, A. Devaraju and S. Harikrishnan, *Mater. Today*, 2018, **5**, 14423–14431.
- 17 B. K. Choure, T. Alam and R. Kumar, *J. Energy Storage*, 2023, **72**, 108161.
- 18 X. Chen, J. Xu, Y. Li, Y. Gao and G. Wang, *SusMat*, 2023, **3**, 510–521.
- 19 K. Zhou, Y. Sheng, W. Guo, L. Wu, H. Wu, X. Hu, Y. Xu, Y. Li, M. Ge, Y. Du, X. Lu and J. Qu, *Adv. Compos. Hybrid Mater.*, 2023, **6**, 34.
- 20 X. Fu, Y. Liu, X. Jiang, Q. Wang, Y. Luo and Y. Lyu, *Thermochim. Acta*, 2018, **665**, 111–118.
- 21 J. Tang, M. Yang, F. Yu, X. Chen, L. Tan and G. Wang, *Appl. Energy*, 2017, **187**, 514–522.
- 22 T. Ren, G. Du, Q. Li, Y. Wang, X. Fu, W. Kong, L. Jiang, J. Lei, P. He and Y. Xiao, *J. Energy Storage*, 2022, **47**, 103592.
- 23 H. Zhang, L. Wang, S. Xi, H. Xie and W. Yu, *Renewable Energy*, 2021, **175**, 307–317.
- 24 X. Zhu, Q. Wang, S. Kang, J. Li and X. Jia, *Chem. Eng. J.*, 2020, **395**, 125112.
- 25 Z. Zou, W. Wu, Y. Wang and D. Drummer, *Mater. Res. Express*, 2020, **7**, 045601.
- 26 Y. Sun, N. Zhang, X. Pan, W. Zhong, B. Qiu, Y. Cai and Y. Yuan, *Int. J. Energy Res.*, 2021, **45**, 20372–20383.
- 27 S. Zhang, Y. Li, Y. Du, X. Ma, J. Lin and S. Chen, *Diam. Relat. Mater.*, 2022, **130**, 109507.
- 28 S. Rawat, T. Boobalan, M. Sathish, S. Hotha and B. Thallada, *Biomass Bioenergy*, 2023, **171**, 106747.
- 29 M. Kamali, L. Appels, E. E. Kwon, T. M. Aminabhavi and R. Dewil, *Chem. Eng. J.*, 2021, **420**, 129946.
- 30 L. Feng, Z. Gao, T. Hu, S. He, Y. Liu, J. Jiang, Q. Zhao and L. Wei, *Chem. Eng. J.*, 2023, **471**, 144772.
- 31 D. Torsello, M. Bartoli, M. Giorcelli, M. Rovere, R. Arrigo, G. Malucelli, A. Tagliaferro and G. Ghigo, *Nanomaterials*, 2021, **11**, 2383.
- 32 C. Liu, M. Han, J. Lin, W. Liu, J. Liu and Z. Zeng, *Carbon*, 2023, **208**, 255–276.
- 33 Y. Zhang, M. He, L. Wang, J. Yan, B. Ma, X. Zhu, Y. S. Ok, V. Mechtcherine and D. C. W. Tsang, *Biochar*, 2022, **4**, 59.
- 34 L. Ke, Q. Wu, N. Zhou, J. Xiong, Q. Yang, L. Zhang, Y. Wang, L. Dai, R. Zou, Y. Liu, R. Ruan and Y. Wang, *Renew. Sustain. Energy Rev.*, 2022, **165**, 112607.
- 35 S. Wang, G. Dai, H. Yang and Z. Luo, *Prog. Energy Combust. Sci.*, 2017, **62**, 33–86.
- 36 E. H. Lee, R. S. Park, H. Kim, S. H. Park, S. C. Jung, J. K. Jeon, S. C. Kim and Y. K. Park, *J. Ind. Eng. Chem.*, 2016, **37**, 18–21.
- 37 Y. Wei, J. Li, F. Sun, J. Wu and L. Zhao, *Green Chem.*, 2018, **20**, 1858–1865.
- 38 Z. Ji, S. Y. H. Abdalkarim, H. Li, R. A. Asad and H. Y. Yu, *Sol. Energy Mater. Sol. Cells*, 2023, **259**, 112453.
- 39 J. He, G. Zhao, P. Mu, H. Wei, Y. Su, H. Sun, Z. Zhu, W. Liang and A. Li, *Sol. Energy Mater. Sol. Cells*, 2019, **201**, 110111.
- 40 S. Liu, S. Peng, B. Zhang, B. Xue, Z. Yang, S. Wang and G. Xu, *RSC Adv.*, 2022, **12**, 9587–9598.
- 41 D. Sarabandi, G. Roudini and F. Barahuie, *J. Energy Storage*, 2019, **24**, 100795.
- 42 J. S. Hua, C. Yuan, X. Zhao, J. Zhang and J. Du, *Energy Sources, Part A*, 2019, **41**, 86–93.
- 43 L. Jiang, Y. Lei, Q. Liu and J. Lei, *Energy*, 2020, **193**, 116802.
- 44 W. Wu, W. Wu and S. Wang, *Appl. Energy*, 2019, **236**, 10–21.
- 45 Z. Shen, S. Kwon, H. L. Lee, M. Toivakka and K. Oh, *Carbohydr. Polym.*, 2021, **273**, 118585.
- 46 L. Liu, X. Zhang and X. Lin, *Renewable Energy*, 2022, **187**, 572–585.
- 47 D. Sarabandi, G. Roudini and F. Barahuie, *J. Energy Storage*, 2019, **24**, 100795.
- 48 T. Khadiran, M. Z. Hussein, Z. Zainal and R. Rusli, *Energy*, 2015, **82**, 468–478.
- 49 S. Paneliya, S. Khanna, A. P. Singh, Y. K. Patel, A. Vanpariya, N. H. Makani, R. Banerjee and I. Mukhopadhyay, *Renewable Energy*, 2021, **167**, 591–599.
- 50 S. Mandal, S. Ishak, D. E. Lee and T. Park, *J. Energy Storage*, 2022, **55**, 105414.
- 51 B. Tan, Z. Huang, Z. Yin, X. Min, Y. G. Liu, X. Wu and M. Fang, *RSC Adv.*, 2016, **6**, 15821–15830.
- 52 Y. Zhao, X. Min, Z. Huang, Y. g. Liu, X. Wu and M. Fang, *Energy Build.*, 2018, **158**, 1049–1062.
- 53 Y. Li, X. Huang, Y. Li, Z. Xi, G. Hai, Z. Tao and G. Wang, *Sustainable Energy Fuels*, 2020, **4**, 1764–1772.
- 54 R. Yang, X. Guo, H. Wu, W. Kang, K. Song, Y. Li, X. Huang and G. Wang, *Biochar*, 2022, **4**, 38.
- 55 L. Lv, S. Huang, C. Zhou and W. Ma, *Ind. Crops Prod.*, 2023, **204**, 117184.
- 56 X. Shi, Y. Meng, R. Bi, Z. Wan, Y. Zhu and O. J. Rojas, *Composites, Part B*, 2022, **245**, 110231.

