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# Thermal conductivity of polyurethane sheets containing beryllium oxide nanofibers

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Polyvinyl alcohol/beryllium sulfate/polyethyleneimine (PVA/BeSO<sub>4</sub>/PEI) precursor nanofibers (NFs) was first fabricated to obtain PVA/BeSO<sub>4</sub>/PEI electrospun NFs by electrospinning technology, finally manufactured beryllium oxide (BeO) NFs followed by various heat treatment methods. The minimum calcination temperature for pure BeO NFs was 1000 °C, and the minimum specific surface area (5.1 m<sup>2</sup> g<sup>-1</sup>) and pore volumes (0.0128 cm<sup>3</sup> g<sup>-1</sup>) were at 1300 °C. 46.18% Be and 53.82% O was measured in BeO NFs by X-ray photoelectron spectroscopy. BeO NFs were then impregnated with polyurethane (PU) aqueous solution to make PU/BeO NFs heat-dissipating sheet. This heat-dissipating sheet showed superior thermal conductivity (14.4 W m<sup>-1</sup> K<sup>-1</sup>) at 41.4 vol% BeO NFs content. The electrical insulating properties of the heat-dissipating sheet were likewise excellent (1.6 × 10<sup>12</sup> Ω □<sup>-1</sup>). In this study, the author attempted to create a thermally conductive but electrically insulating PU/BeO NFs heat-dissipating sheet that could effectively eliminate generated heat from electric equipment.

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## Introduction

Electronic devices and other high-performance energy-saving home appliances are now significantly smaller, with higher integration and power. Continuous using such devices generates a large amount of heat. The electronic components may create various problems, such as increased failure frequency, fire, and smoke, when the increased temperature exceeds the thermal stability limit of this device. Many researchers tried to develop heat-dissipating materials to improve the thermal conductivity of electronic devices. Some researchers used metal as heat-dissipating materials for electronics devices because they possess excellent thermal conductivity and superior mechanical strength. Although, electrically insulating properties are vital for developing electric devices.<sup>1–4</sup>

Polymeric materials possess excellent electric insulation properties, good moldability, and thermal insulating properties. So, polymeric materials can be used as heat-dissipating sheets if the thermal conductivity properties improve. Adding a certain amount of inorganic filler with low thermal conductive polymer (0.01–0.1 W m<sup>-1</sup> K<sup>-1</sup>), the thermal conductivity of polymeric composite will increase significantly.<sup>5,6</sup> The most significant advantages of polymer composites are their simple processing technique, very lightweight, meager cost, and resistance against corrosions.<sup>7–13</sup> Some inorganic materials possess

thermal conductivity with electrical insulation properties, such as silicon nitride (Si<sub>3</sub>N<sub>4</sub>),<sup>14,15</sup> beryllium oxide (BeO),<sup>16,17</sup> hexagonal boron nitride (h-BN),<sup>18–20</sup> aluminum nitride (AlN),<sup>21,22</sup> alumina (Al<sub>2</sub>O<sub>3</sub>)<sup>23</sup> and silicon carbide (SiC).<sup>24</sup>

One of the most critical properties of BeO ceramic is its ultra-high thermal conductivity (330 W m<sup>-1</sup> K<sup>-1</sup>) with excellent electrical insulation (>10<sup>13</sup> Ω cm) properties, among other well-known ceramic materials. However, depending on density, the thermal conductivity of BeO ceramic varies from 230 to 330 W m<sup>-1</sup> K<sup>-1</sup>, which is higher than other metals except for copper, silver, and gold.<sup>17</sup> Other than high thermal conductive properties, beryllium ceramics can exhibit a unique combination of other physicochemical properties such as high melting point, high intensity, high insulation nature, high chemical and thermal stability, a considerable specific volumetric resistance, transparency for vacuum, infrared, ultraviolet, visible, radiation resistance, low dielectric losses, low dielectric constant, good technology applicability. BeO ceramics can conduct heat from electronic technology while temperatures range from 300 to 630 K of all ceramic materials. This outstanding property makes BeO ceramic a promising material for using new technology fields, contemporary electronics, vacuum electronics technology, nuclear technology, and microelectronics. There is no alternative to BeO ceramics, especially in thermal applications.<sup>25,26</sup>

There are few reports on BeO nanoparticle synthesis, and extensive structural characterization of produced BeO nanoparticles is limited and not reported.<sup>27</sup> Wang *et al.* prepared BeO nanoparticles by polyacrylamide gel route. They created xerogel by using acrylamide and *N,N'*-methylene bis acrylamide monomers and mixed with beryllium sulfate tetrahydrate

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( $\text{BeSO}_4 \cdot 4\text{H}_2\text{O}$ ) aqueous solution followed by ammonium persulfate aqueous solution. Finally, they prepared BeO nanoparticles in white powder by heating and calcinating at 700–1000 °C.<sup>26</sup>

NF research, development, and industrial applications have received much attention recently. Many investigations on heat-dissipating sheets constructed of NFs have been undertaken.<sup>28,29</sup> Ohgoshi *et al.* fabricated magnesia (MgO) NFs by calcinating polyvinyl alcohol/magnesium ethoxide ( $\text{PVA}/\text{Mg}(\text{OC}_2\text{H}_5)_2$ ) precursor NFs at different temperatures. They produced  $\text{PVA}/\text{Mg}(\text{OC}_2\text{H}_5)_2$  precursor NFs mat by electrospinning technique using PVA and  $\text{Mg}(\text{OC}_2\text{H}_5)_2$  mixtures.<sup>30</sup> Nakane *et al.* created  $\alpha$ -alumina NF mats from a PVA/boehmite spinning mix using electrospinning. They discovered that at 5.7%  $\alpha$ -alumina content, the  $\alpha$ -alumina NFs sheet had thermal conductivity in parallel ( $1.29 \text{ W m}^{-1} \text{ K}^{-1}$ ) and perpendicular ( $0.38 \text{ W m}^{-1} \text{ K}^{-1}$ ) directions.<sup>31,32</sup> Ohgoshi *et al.* generated an  $\alpha$ -alumina NFs polyurethane sheet with boehmite particles and aqueous PVA solution using an electrospinning method. The resultant  $\alpha$ -alumina NFs polyurethane composite sheet had increased heat conductivity ( $19.7 \text{ W m}^{-1} \text{ K}^{-1}$ ) in a parallel direction when it included 33% alumina.<sup>33,34</sup>

Electrospinning is becoming a popular technology for polymer NFs since it is simple to make various inorganic NFs. Electrospinning creates continuous polymer strands with nanoscale diameters by applying an external electric field to a polymer solution.<sup>35</sup> Compared to commercial textiles, electrospun NFs have a large specific surface area and a tiny pore size. As a result, NFs can be rationally designed to have novel and significantly improved physical, chemical, and biological properties.<sup>36</sup> An electric force is applied between a suspended droplet solution at a capillary tip and a collector in the basic process of electrospinning. A charged jet is released and travels to the grounded target when the intensity of the electric field overcomes the surface tension of the polymer solution, creating fibers in the form of nonwoven mats.<sup>37</sup>

The authors used granular polyvinyl alcohol (PVA) and  $\text{BeSO}_4 \cdot 4\text{H}_2\text{O}$  salt and polyethyleneimine (PEI) to produce PVA/BeO/PEI precursor NFs in this research work. After impregnating polyurethane (PU), they make PU/BeO NFs heat-dissipating sheets from PVA/BeO/PEI precursor NFs. PU/BeO NFs heat-dissipating sheet may be very effective as a heat sink because BeO ceramic has high thermal conductivity and excellent electrical resistivity. On the other side, this study also focuses on the electrospinning technique to make BeO NFs, followed by the heat treatment method and make PU/BeO NFs composite sheet. This composite sheet is most effective in dissipating heat from electronic devices. So BeO NFs may be very effective for high-power electronic devices dissipating heat.

## Experimental section

### Raw materials

$\text{BeSO}_4 \cdot 4\text{H}_2\text{O}$  (purity > 98%) was purchased from Kanto chemical co. Inc., Tokyo, Japan. PVA, Degree of polymerization = 1500 and molecular weight = 67 000 was purchased from Wako Pure Chemical Ind. Ltd., Japan. PEI ( $\text{CH}_2\text{CH}_2\text{NH}$ )<sub>n</sub> (30% in

water with a molecular weight of 60 000–80 000) was purchased from Nacalai tesque, Inc., Kyoto, Japan. Deionized water ( $\text{H}_2\text{O}$ ) was used as received. PU water emulsion containing 30 wt%, solid (Superflex #300), a kind-hearted gifted by DKS Co. Ltd., Japan. All chemicals were used without any further purification.

### BeO NFs preparation by electrospinning machine

10 g of PVA were added with 90 g of deionized water and boiled for 4 hours at 90 °C to make a 10% PVA solution. To make the PVA/ $\text{BeSO}_4$  solution,  $\text{BeSO}_4 \cdot 4\text{H}_2\text{O}$  salt was added directly to the previously produced 10% PVA solution and kept at room temperature for 7 hours with vigorous stirring. PEI was added with PVA/ $\text{BeSO}_4$  solution, finally making PVA/ $\text{BeSO}_4$ /PEI composite precursor solution. The electrospinning apparatus was used to make PVA/ $\text{BeSO}_4$ /PEI precursor NFs. The prepared precursor solutions were placed in a 3 mL plastic syringe with a 0.5 mm pinhead inner diameter. The electrospinning settings in this experiment were as follows: 20 kV operating voltage, 0.008 mL min<sup>-1</sup> injection rate, and 15 cm needle tip to collector distance. The electrospun PVA/ $\text{BeSO}_4$ /PEI composite NFs were collected on parchment paper. PVA/ $\text{BeSO}_4$ /PEI precursor NFs were calcined at varied heat treatment temperatures (700 °C, 800 °C, 900 °C, 1000 °C, 1100 °C, and 1200 °C) for 5 hours in the air with an electric furnace to eliminate organic components (NHK-170, Nitto Kagaku Co. Ltd., box-type furnace, Japan). Finally, the BeO NFs were then obtained.

### Formation of PU sheets containing BeO NFs

The BeO NFs were impregnated with a 5% PU solution and cured for 10 hours by vacuuming the solution at room temperature until completely absorbed. Fig. 1 depicted all of the appropriate steps schematically.

### Characterization

The morphological structure of both precursor and BeO NFs was examined at a voltage of 5 kV using a Keyence scanning electron microscope (VE-9800, Keyence Co. Ltd., Japan). The samples were initially coated with Au/Pd sputtering under vacuum using an ion coater (SC-701; Sanyu Electron Co. Ltd., Japan). Adobe Photoshop CS3 extension software measurements of the precursor and BeO NFs ( $n = 100$ ) were used to calculate the mean fiber diameter  $D$  and standard deviation. An X-ray diffractometer (XRD, Rigaku Mini Flex II, Japan) was used to measure X-ray diffraction using Ni-filtered CuK radiation (30 kV, 15 mA) as an X-ray source to investigate the samples' crystalline nature. The BeO content (wt%) in the dissipation sheets was assessed by thermogravimetric (TG) analysis carried out in the air atmosphere with a heating speed of 5 °C min<sup>-1</sup> from 30 to 600 °C using a differential TG analyzer (Shimadzu DTG-60, Japan). A BELSORP-mini II (MicrotracBEL Corp., Japan) was used to monitor nitrogen adsorption isotherms at -196 °C. Specific surface areas were calculated using the Brunauer–Emmett–Teller (BET) method. The Barrett–Joyner–Halenda (BJH) algorithm was used to calculate the pore-size distribution (PSD) curves from the isotherm. A JEOL JPS-9010 (Nippon Electronics Co. Ltd., Japan) equipment was used to perform X-



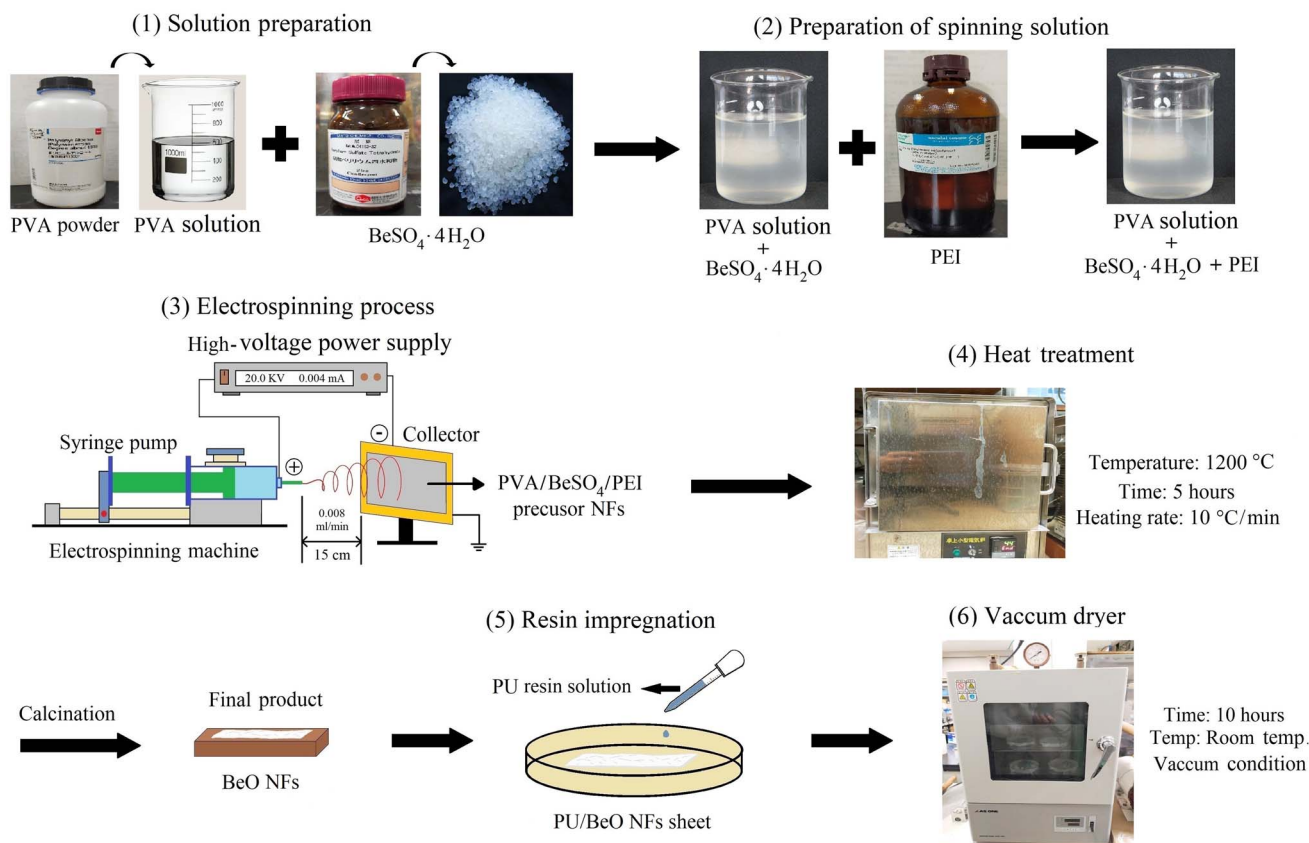


Fig. 1 Experimental scheme for the preparation of BeO NF mats.

ray photoelectron spectroscopy (XPS) using an Mg-Kα ( $h\nu = 1253.6$  eV) source at a residual gas pressure of  $5 \times 10^{-6}$  Pa. Hiresta-UP (Mitsubishi Chemical Analytics, Japan) was used to measure the electrical resistivity of the composite sheet by ASTM D257. Thermal diffusivity ( $\text{m}^2 \text{s}^{-1}$ ) was measured at room temperature in the planar and thickness directions using a thermowave analyzer TA35 (Bethel Co., Ltd., Japan). Thermal diffusivity is measured *via* temperature wave analysis in this system. We determined four different average and relative standard deviation (RSD) measurements.

## Results and discussion

### Characterization of BeO NFs

In our previous work, we heated PVA/BeSO<sub>4</sub>/PEI precursor NFs' from 700 °C to 1200 °C.<sup>38</sup> However, here we start heating from 600 °C to 1300 °C. The XRD curves of the residues calcinated from 600 °C to 1300 °C are shown in Fig. 2. Few low-intensity BeO crystal peaks are observed after heating from 600 °C to 900 °C. However, the most pronounced BeO crystal peaks were detected at temperatures of 1000 °C or higher, and their intensity increased as the temperature grew. The main reason is that all organic components are thermally decomposed at high temperatures (1000 °C to 1300 °C). However, BeO residues is very brittle while heated at 1200 °C and above.

The calcination temperatures can influence the surface area and morphologies of fibers, tuning the crystalline structure.

PVA/BeSO<sub>4</sub>/PEI precursor NFs' specific surface area ( $36.3 \text{ m}^2 \text{g}^{-1}$ ) increases significantly to  $145 \text{ m}^2 \text{g}^{-1}$  for BeO NFs while calcinated at 600 °C. Furthermore, within the mesopore range of 2–50 nm, a mesoporous structure with an average pore diameter of 9.3 nm is seen. However, as the calcination

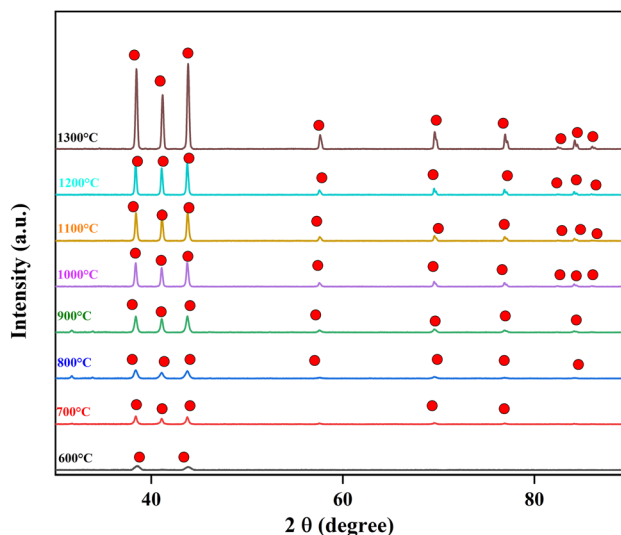


Fig. 2 XRD curves of residues after calcinating PVA/BeSO<sub>4</sub>/PEI precursor NFs (●: BeO crystal).

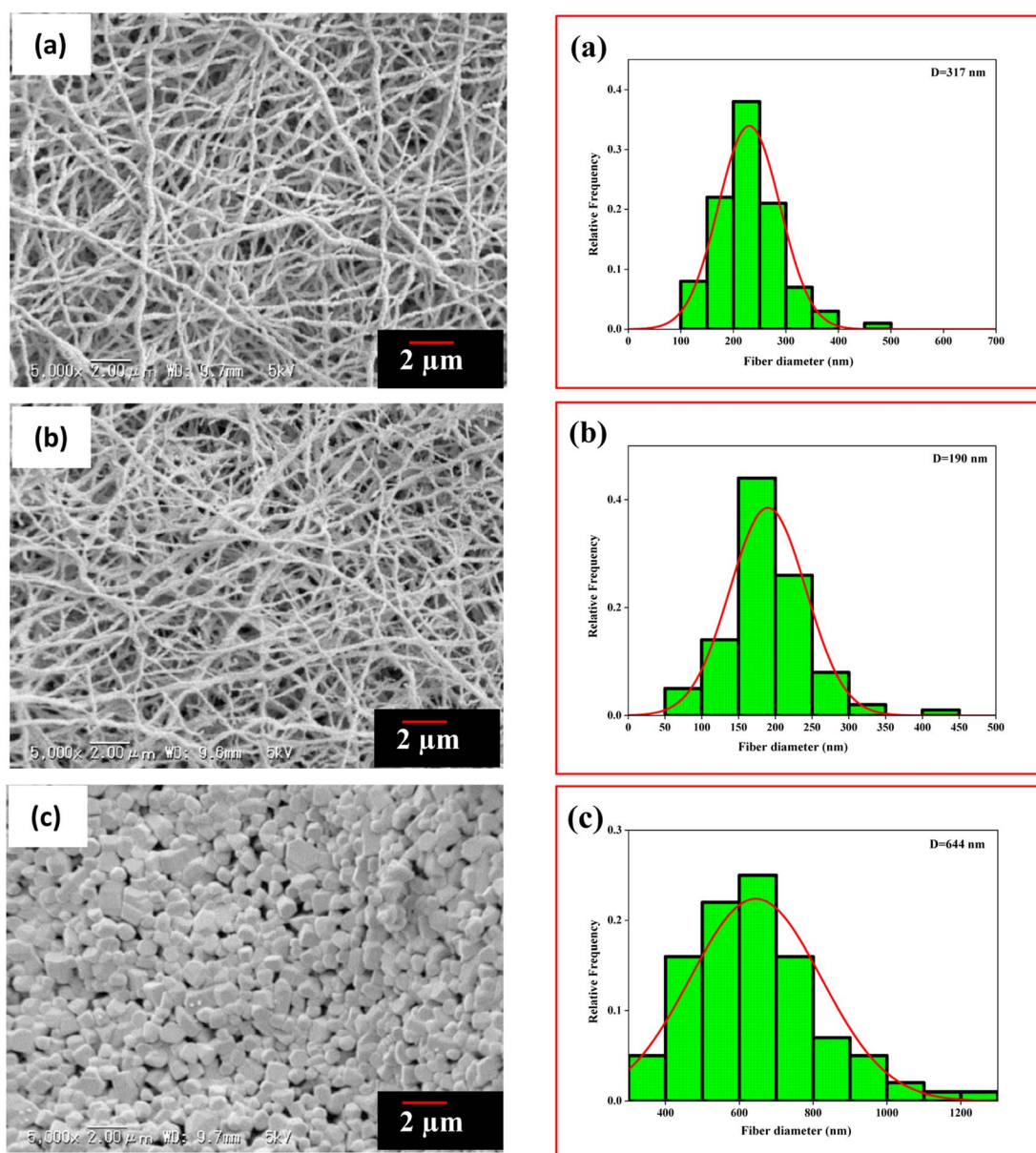




**Table 1** Pore characteristics of each residue at different heating temperatures

Precursor and calcinated samples	Specific surface area ( $\text{m}^2 \text{g}^{-1}$ )	Pore volume ( $\text{cm}^3 \text{g}^{-1}$ )
PVA/ $\text{BeSO}_4$ /PEI precursor NFs	36.9	0.0638
BeO NFs at 600 °C	145.0	0.3393
BeO NFs at 700 °C	122.0	0.4318
BeO NFs at 800 °C	77.4	0.2891
BeO NFs at 900 °C	50.8	0.1827
BeO NFs at 1000 °C	23.0	0.0643
BeO NFs at 1100 °C	17.0	0.0382
BeO NFs at 1200 °C	11.9	0.0275
BeO residues at 1300 °C	5.1	0.0128

temperature rises to 1000 °C, 1100 °C, 1200 °C, and 1300 °C, the specific surface area reduces dramatically to  $23.0 \text{ m}^2 \text{g}^{-1}$ ,  $17.0 \text{ m}^2 \text{g}^{-1}$ ,  $11.9 \text{ m}^2 \text{g}^{-1}$ , and  $5.1 \text{ m}^2 \text{g}^{-1}$ , respectively. The findings show that removing organics improves specific surface area at an adequate calcination temperature. A high calcination temperature will hinder the increase in specific surface area, resulting in a low specific surface area. Sintering or metal particle obstruction could be the cause. At temperatures of 1000 °C or higher, BeO NFs can be employed as heat-dissipating sheets. However, when heat-treated at 1200 °C and 1300 °C temperatures, samples containing BeO crystal became brittle, as the fiber shape was shattered. Based on nitrogen adsorption investigations, Table 1 summarizes the specific surface area, average pore diameter, and pore volume estimates.

**Fig. 3** SEM images and histogram of residues after calcinating PVA/ $\text{BeSO}_4$ /PEI precursor NFs at (a) 600 °C, (b) 1100 °C, (c) 1300 °C.

The electrospinning method fabricated the PVA/BeSO<sub>4</sub>/PEI precursor NFs. On the other hand, BeO crystals were formed by calcinating the PVA/BeSO<sub>4</sub>/PEI precursor NFs at 1000 °C or above temperatures at a dry ratio of PVA/BeO was 90/10 wt%;<sup>38</sup> the NFs remained in their shape, and the average fiber diameter was reduced due to heat treatment up to 1200 °C temperatures.<sup>39</sup> SEM images and histogram of BeO residue after calcinating PVA/BeSO<sub>4</sub>/PEI precursor NFs at 600 °C, 1100 °C, and 1300 °C temperatures are shown in Fig. 3. At 600 °C and 1100 °C temperatures, BeO remaining good NFs shape, interestingly, BeO crystal became very compact and cubic in shape at 1300 °C temperatures, as shown in Fig. 3(c).

The ATR-FTIR spectra of pure PVA, BeSO<sub>4</sub>·4H<sub>2</sub>O salt, PVA/BeSO<sub>4</sub>/PEI precursor NFs, and BeO NFs were obtained, and the results are shown in Fig. 4. The spectra show strong broadband for pure PVA at 3320 cm<sup>-1</sup> and 3210 cm<sup>-1</sup> for PVA/BeSO<sub>4</sub>/PEI precursor NFs. This band was attributed to the O–H stretching vibration of the hydroxyl group of pure PVA and PVA/BeSO<sub>4</sub>/PEI precursor NF samples because water contains hydrogen bonding. However, no absorbance peak was found for the BeO NFs due to a lack of water-containing hydrogen bonding for BeO NF samples.<sup>40</sup> The band corresponding to C–H stretching mode is 2926 cm<sup>-1</sup> and 2940 cm<sup>-1</sup> for the pure PVA and PVA/BeSO<sub>4</sub>/PEI precursor NFs.<sup>41</sup> The absorption peak at 853 cm<sup>-1</sup> for pure PVA has been assigned to C–H rocking (Fig. 4a) and moved to 823 cm<sup>-1</sup> for PVA/BeSO<sub>4</sub>/PEI precursor NFs (Fig. 4c). The sharp band of 1094 cm<sup>-1</sup> corresponds to the C–O stretching of acetyl groups present on the PVA backbone that is shifted to 1060 cm<sup>-1</sup> and 1080 cm<sup>-1</sup> for BeSO<sub>4</sub>·4H<sub>2</sub>O salt and PVA/BeSO<sub>4</sub>/PEI precursor NFs, respectively.<sup>40–42</sup> In the complexed system, CH<sub>2</sub> bending exhibited at 1426 cm<sup>-1</sup> in pure PVA remains the same for PVA/BeSO<sub>4</sub>/PEI precursor NFs. The acetyl C=O group has an absorbance of 1727 cm<sup>-1</sup> in pure PVA, which can be explained by intra/inter molecule hydrogen bonding with the neighboring OH group. For instance, this C=O peak is shifted

to 1651 cm<sup>-1</sup> and 1734 cm<sup>-1</sup> for BeSO<sub>4</sub>·4H<sub>2</sub>O salt and PVA/BeSO<sub>4</sub>/PEI precursor NFs, respectively. This demonstrates that the interaction between the salt and PVA is caused by pure PVA's O–H and C–O groups. Water molecules are likely to be removed at higher calcinating temperatures (1100 °C) could be the reason.<sup>40,41</sup> Furthermore, the BeO bending vibration was measured below 1000 cm<sup>-1</sup>, similar to other inorganic metal oxide compounds.<sup>43–48</sup> The low peak is  $\nu$  (Be–O) vibrations at 725 cm<sup>-1</sup>.<sup>49</sup>

BeO NFs were subjected to XPS examination to determine which chemical elements were present on the surface of these compounds. In contrast, BeSO<sub>4</sub>·4H<sub>2</sub>O salt, PVA/BeSO<sub>4</sub> precursor NFs and PVA/BeSO<sub>4</sub>/PEI precursor NFs were used as controls. In Fig. 5, BeSO<sub>4</sub>·4H<sub>2</sub>O salt, PVA/BeSO<sub>4</sub>, and PVA/BeSO<sub>4</sub>/PEI precursor NFs showed peaks of O 1s, C 1s, and S 2p<sub>3/2</sub>. However, the BeO NFs showed pronounced O 1s peaks where C 1s and S 2p<sub>3/2</sub> peaks were absent. Be 1s peaks were present only in the BeO NFs but not in the BeSO<sub>4</sub>·4H<sub>2</sub>O salt, PVA/BeSO<sub>4</sub> precursor NFs, or PVA/BeSO<sub>4</sub>/PEI precursor NFs.

Fig. 6 shows the BeSO<sub>4</sub>·4H<sub>2</sub>O salt, PVA/BeSO<sub>4</sub> precursor NFs, PVA/BeSO<sub>4</sub>/PEI precursor NFs and BeO NFs samples C 1s, O 1s, Be 1s, and S 2p<sub>3/2</sub> XP spectra. The PVA/BeSO<sub>4</sub>/PEI precursor NFs sample spectra showed strong C–C bonding at 284.2 eV (Fig. 6a). Compared with the BeO NF samples, the C 1s peak increased in the PVA/BeSO<sub>4</sub>/PEI precursor NF samples. The main reason is the significant amount of carbon in precursor samples. The O 1s peaks were detected in all instances; however, the sharpest rise (531.7 eV) for BeO NFs (Fig. 6b). The main reason is that a higher calcinating temperature (1100 °C) reduces organic components. On the other hand, Be 1s peaks were detected at a very high calcinating temperature (1100 °C), only for BeO NFs at 113.7 eV (Fig. 6c). In addition, S 2p<sub>3/2</sub> peaks were absent in all samples (PVA/BeSO<sub>4</sub> precursor NFs, PVA/BeSO<sub>4</sub>/PEI precursor NFs and BeO NFs) except BeSO<sub>4</sub>·4H<sub>2</sub>O salt (Fig. 6d) at 168.1 eV.

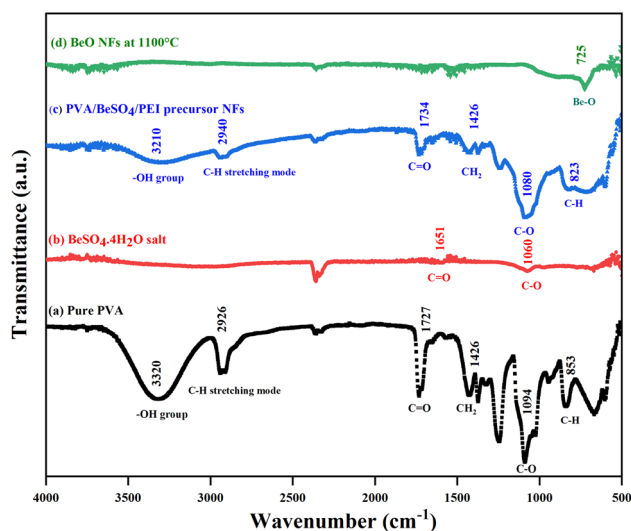


Fig. 4 FTIR spectra of (a) pure PVA, (b) BeSO<sub>4</sub>·4H<sub>2</sub>O salt, (c) PVA/BeSO<sub>4</sub>/PEI precursor NFs, and (d) BeO NFs.

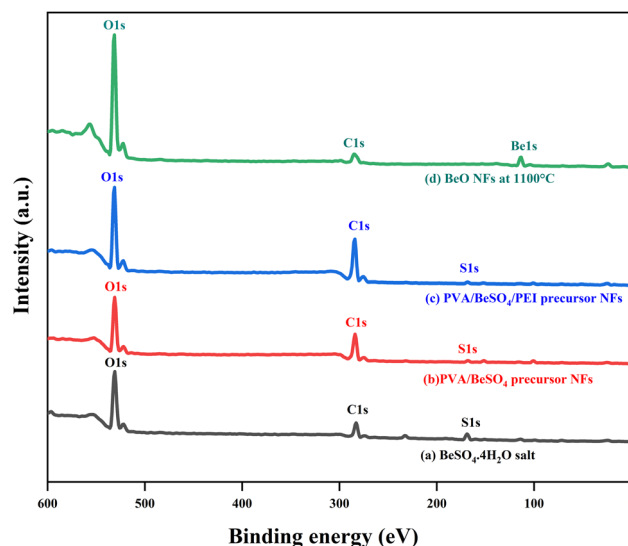


Fig. 5 Wide-scan XPS spectra of BeSO<sub>4</sub>·4H<sub>2</sub>O salt, PVA/BeSO<sub>4</sub> precursor NFs, PVA/BeSO<sub>4</sub>/PEI precursor NFs and BeO NFs.



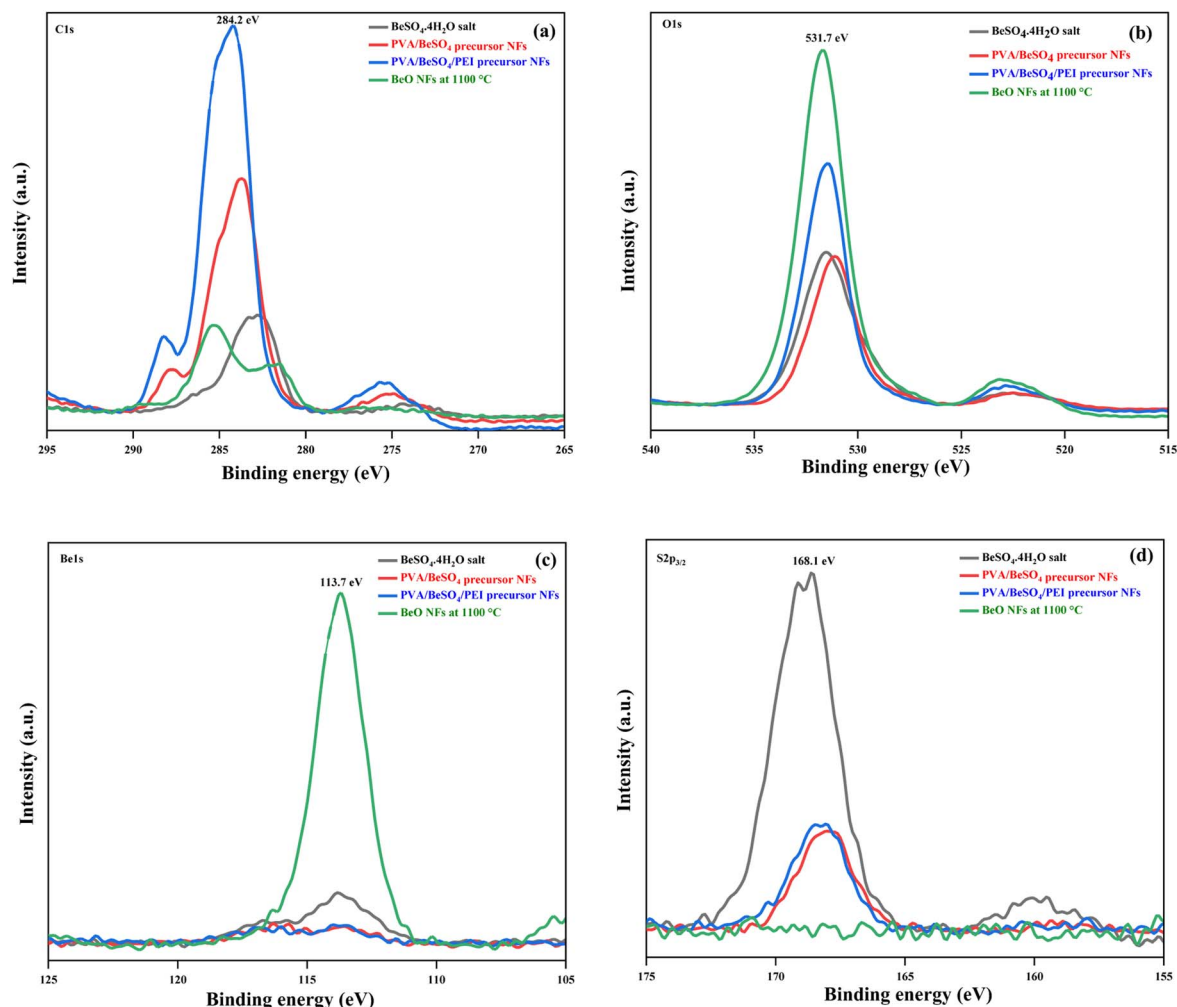


Fig. 6 XPS spectra of  $\text{BeSO}_4 \cdot 4\text{H}_2\text{O}$  salt, PVA/ $\text{BeSO}_4$  precursor NFs, PVA/ $\text{BeSO}_4$ /PEI precursor NFs and BeO NFs (a) C 1s spectra, (b) O 1s spectra, (c) Be 1s spectra, and (d) S  $2p_{3/2}$  spectra.

The elemental composition contents (Table 2) of C, O, Be, S, and N were analyzed according to the XPS results (Fig. 5) to corroborate the variations in the temporal composition content of each sample. Table 2 shows that no elements other than Be (46.18%) and O (53.82%) could be detected. Be's weight percentage was significantly increased at BeO NFs compared to  $\text{BeSO}_4 \cdot 4\text{H}_2\text{O}$  salt and precursor NFs. The possible reason is that the Be content increased dramatically with decreasing C content after calcination, which may lead to BeO bonds.

Table 2 Elemental composition contents of  $\text{BeSO}_4 \cdot 4\text{H}_2\text{O}$  salt, PVA/ $\text{BeSO}_4$  precursor NFs, PVA/ $\text{BeSO}_4$ /PEI precursor NFs and BeO NFs evaluated from XPS results

Sample name	Elemental composition contents (%)				
	C 1s	Be 1s	O 1s	S $2p_{3/2}$	N 1s
$\text{BeSO}_4 \cdot 4\text{H}_2\text{O}$ salt	24.75	18.83	48.97	7.45	—
PVA/ $\text{BeSO}_4$ precursor NFs	53.28	7.94	37.11	1.67	—
PVA/ $\text{BeSO}_4$ /PEI precursor NFs	57.07	5.14	36.42	1.20	0.17
BeO NFs heated at 1100 °C	—	46.18	53.82	—	—

### Formation of PU/BeO NFs composite sheet

SEM images of the PU sheet containing the BeO NFs are shown in Fig. 7. PU is impregnated into the calcinated BeO NF mats, making the sheet significantly closer. Here, the PU successfully filled the gap between the NFs, and the NFs were dispersed homogeneously in the PU matrix. NFs were also found on the sheet's surface, indicating that the fibers were neither fractured nor detached once soaked with PU emulsion. However, there are very few heat transmission channels for BeO in the thickness direction of the sheet. We assumed that the produced sheets would have a superior thermal conductivity in the plane direction than in the thickness direction.

### Thermal conductivity of PU/BeO NF composite sheet

The brittleness of the as-prepared BeO NF mats prevented measuring their thermal conductivity. However, the thermal conductivity of PU/BeO NF composite sheets were measured as shown in Table 3. It was observed that the thermal conductivity was much higher while calcinating at 1100 °C for both plane and thickness directions than at low calcinating temperatures





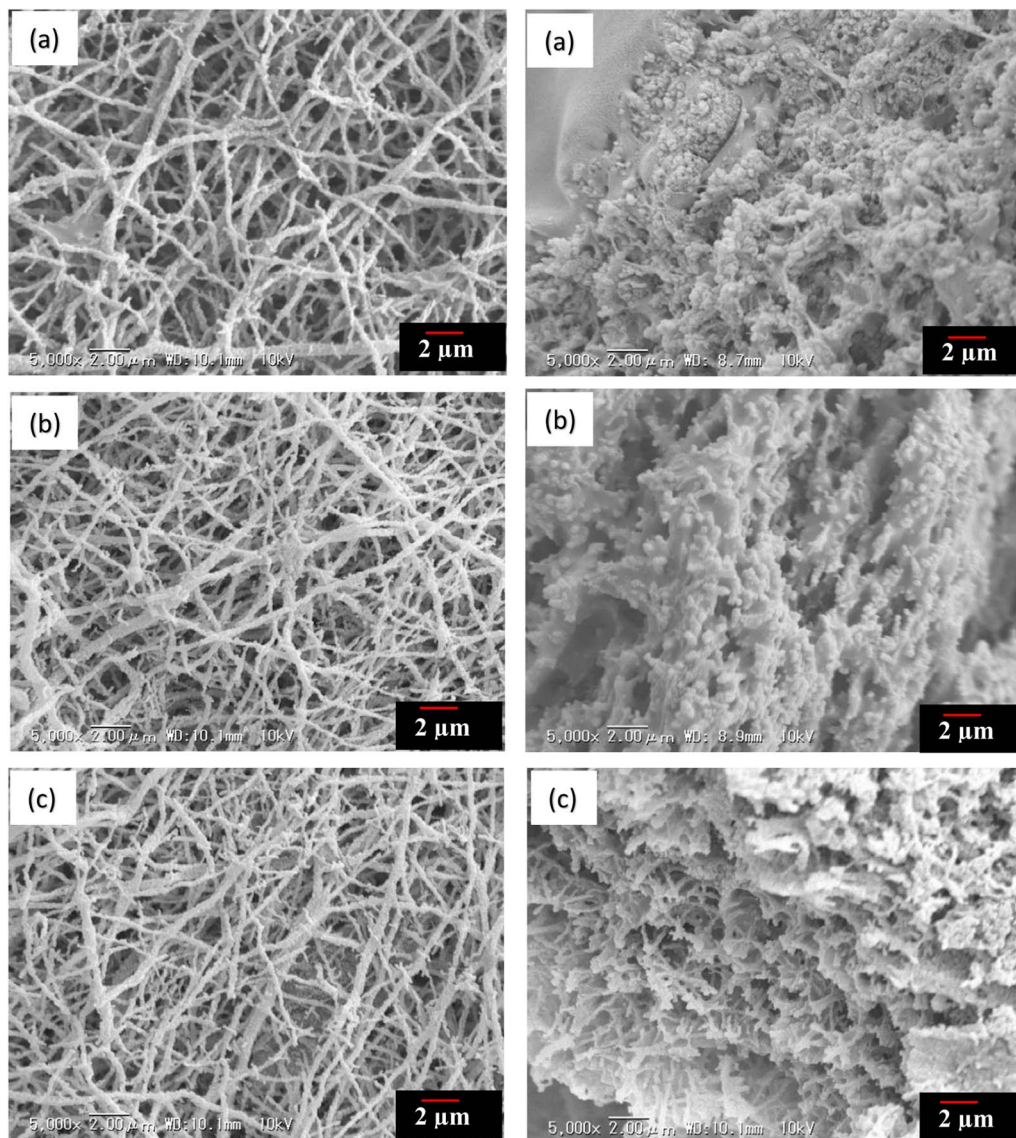


Fig. 7 SEM images of PU sheet containing aligned BeO NFs (a: 700 °C, b: 900 °C, c: 1100 °C, (left: sheet surface, right: cross-section)).

**Table 3** Thermal conductivity of BeO heat-dissipating sheet at 700 °C, 900 °C, and 1100 °C calcination temperatures both in plane and thickness direction

Temperature (°C)	BeO content (%)	Thermal conductivity ( $\text{W m}^{-1} \text{K}^{-1}$ )	
		Plane direction	Thickness direction
700	31.2	$3.0 \pm 0.17^a$	$1.2 \pm 0.75^e$
	40.5	$2.0 \pm 0.20^a$	$0.6 \pm 0.62^f$
	51.1	$4.0 \pm 0.10^a$	$0.5 \pm 0.26^g$
900	21.8	$3.1 \pm 0.43^b$	$2.6 \pm 0.78^h$
	27.5	$3.8 \pm 0.44^c$	$2.3 \pm 1.35^a$
	29.8	$4.0 \pm 0.46^a$	$2.5 \pm 1.32^g$
1100	16.6	$1.4 \pm 0.36^d$	$2.6 \pm 0.65^h$
	21.5	$5.0 \pm 1.39^c$	$4.4 \pm 1.68^i$
	22.9	$4.5 \pm 0.62^a$	$0.8 \pm 0.53^b$
	41.4	$14.4 \pm 0.78^a$	$1.4 \pm 0.56^e$

(700 °C and 900 °C). The possible reason is that due to high calcination temperatures, organic content removes, and BeO NFs behave as mesoporous NFs.

For all the listed data, the standard deviations of thermal conductivity at the plane direction were less than 1.39, and the thickness direction was less than 1.68, which was presented as percentages of  $L_s$  (mean  $\pm$  SD). To measure the statistical significance, a two-tailed *t*-test was conducted between the datasets, and *p* values were derived, where  $p > 0.05$  indicates no significant difference and  $p \leq 0.05$  indicates a significant difference between the datasets. Here, the statistical analysis showed that, in most cases, the datasets are significantly different ( $p \leq 0.05$ ). So, it can be said that changing calcination temperatures significantly affected ( $p \leq 0.05$ ) the thermal conductivity of PU/BeO NF sheets. Superscripts indicate the test results of the ANOVA. Different superscripts showing datasets are significantly different, and similar superscripts showing



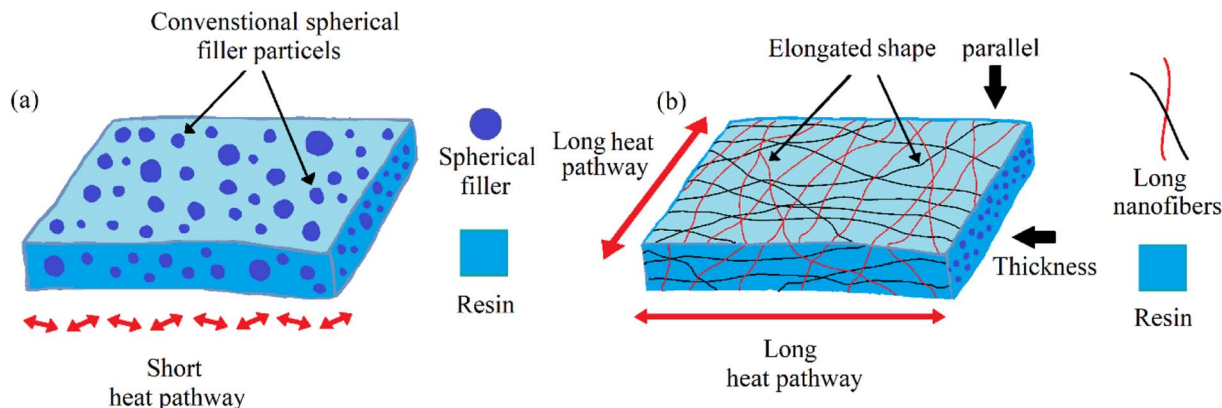


Fig. 8 Mechanism of (a) low thermal conductivity by conventional spherical fillers and (b) improved thermal conductivity by the elongated shape of BeO NFs.

datasets are similar. Here total nine (9) superscripts has been measured ( $a = 0.00$ ,  $b = 0.12$ ,  $c = 0.19$ ,  $d = 0.16$ ,  $e = 0.11$ ,  $f = 0.31$ ,  $g = 0.26$ ,  $h = 0.01$ , and  $i = 0.02$ ).

It also observed that thermal conductivity was much higher in the planner direction than in the thickness direction. The possible reason is that the thermal conductivity in the thickness direction is almost similar to conventional spherical filler particles. The contact area of traditional spherical fillers is limited, and an efficient heat conduction path is difficult to implement. On the other side, the thermal conductivity parallel to the aligned BeO NFs rose in proportion to the NF content. This improvement was due to the BeO NFs' elongated shape, which provides excellent heat pathways for the BeO NFs in the resin. The whole mechanism of low and high thermal conductivity is shown in Fig. 8.

The thermal conductivity of pure is PU,  $0.022 \text{ W m}^{-1} \text{ K}^{-1}$ .<sup>31</sup> Adding  $\text{BeSO}_4 \cdot 4\text{H}_2\text{O}$  salt with pure PU increases the thermal conductivity of the PU/BeO NF sheets. The lowest thermal conductivity ( $1.4 \text{ W m}^{-1} \text{ K}^{-1}$ ) of the PU/BeO NFs sheet was at

16.6 vol% BeO content, which was increased at 21.5, 22.9, 23.4 vol% BeO content. At 41.4 vol% BeO content, the highest thermal conductivity ( $14.4 \text{ W m}^{-1} \text{ K}^{-1}$ ) was found in the plane direction. However, the relationship between BeO% and thermal conductivity is not always regular. The thickness of the highest thermal conductive ( $14.4 \text{ W m}^{-1} \text{ K}^{-1}$ ) BeO NF sheet was  $81 \mu\text{m}$  at 41.4 vol% and content. Fig. 9 shows the thermal conductivity of the PU/BeO NF sheet at  $1100^\circ\text{C}$  temperatures in both plane and thickness directions.

The authors also measured the thermal diffusivity of the PU/BeO NF sheets to predict cooling processes and to know how quickly a material reacts to a change in temperature. Fig. 10 shows the thermal diffusivity of PU/BeO NF sheets at different temperatures. Here, the thermal diffusivity was lower ( $7.7 \times 10^{-7}$  to  $16.5 \times 10^{-7} \text{ m}^2 \text{ s}^{-1}$ ) at  $700^\circ\text{C}$  temperatures compared to  $900^\circ\text{C}$  temperatures ( $12.9 \times 10^{-7}$  to  $16.7 \times 10^{-7} \text{ m}^2 \text{ s}^{-1}$ ). However, the thermal diffusivity was much higher ( $10.8 \times 10^{-7}$  to  $56.7 \times 10^{-7} \text{ m}^2 \text{ s}^{-1}$ ) at higher  $1100^\circ\text{C}$  temperatures than at low temperatures ( $700^\circ\text{C}$  and  $900^\circ\text{C}$ ).

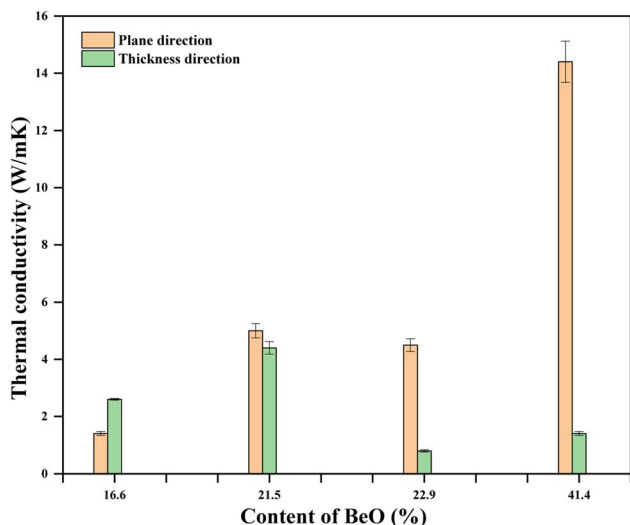


Fig. 9 Effect of BeO content of PU/BeO NF sheets on the thermal conductivity at  $1100^\circ\text{C}$ .

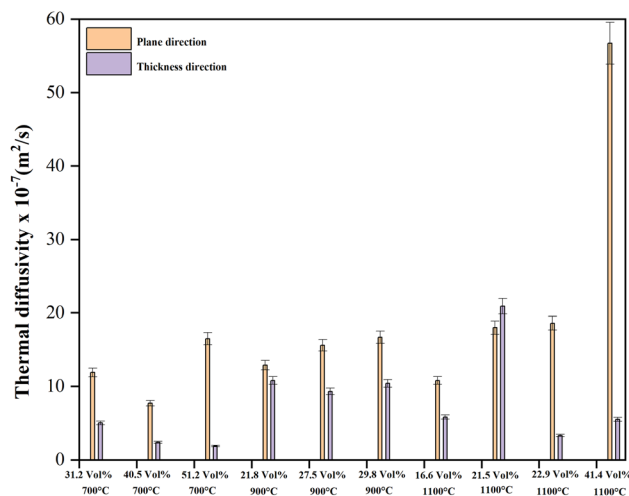


Fig. 10 Thermal diffusivity of PU/BeO NF sheets at  $700^\circ\text{C}$ ,  $900^\circ\text{C}$ , and  $1100^\circ\text{C}$ .





Table 4 Electrical insulation properties of PU/BeO NF sheets

BeO content (%)	Thickness ( $\mu\text{m}$ )	Electrical insulation ( $\Omega \square^{-1}$ )		
		700 $^{\circ}\text{C}$	900 $^{\circ}\text{C}$	1100 $^{\circ}\text{C}$
31.2	132	$1.6 \times 10^{12}$	—	—
40.5	101	$2.4 \times 10^{12}$	—	—
80.4	61	$1.9 \times 10^{12}$	—	—
21.8	100	—	$2.6 \times 10^{12}$	—
27.5	33	—	$2.1 \times 10^{12}$	—
39.0	68	—	$1.6 \times 10^{12}$	—
16.6	552	—	—	$1.8 \times 10^{12}$
21.5	107	—	—	$2.1 \times 10^{12}$
22.9	555	—	—	$1.8 \times 10^{12}$
23.4	155	—	—	$1.5 \times 10^{12}$
41.4	81	—	—	$1.6 \times 10^{12}$

Our previous work could produce a thermal conductivity ( $7.9\text{--}14.4 \text{ W m}^{-1} \text{ K}^{-1}$ ) PU/ $\text{Al}_2\text{O}_3$  NFs heat-dissipating sheet.<sup>39</sup> At this time, we can also create PU/BeO NFs heat-dissipating sheet with high thermal conductivity ( $1.4\text{--}14.4 \text{ W m}^{-1} \text{ K}^{-1}$ ). However, we may be able to improve the thermal conductivity of BeO NFs by overcoming two difficulties. The first difficulty is that we couldn't produce completely nonporous BeO NFs.<sup>39</sup> Another is that we couldn't create high crystallinity of BeO NFs because, after 1200  $^{\circ}\text{C}$  temperatures, BeO crystal is very brittleness. We may circumvent those two challenges by altering the precursor-making process. In addition, the authors strongly believe that mechanical properties are significant from the application point of view. At this time, the authors couldn't create high crystallinity of BeO NF because, after 1200  $^{\circ}\text{C}$  temperatures, BeO crystal is very brittle. In the future, the authors will try to modify the precursors-making process to produce high crystallinity of BeO NF at low temperatures.

#### Electrical insulation properties of BeO/PU sheets (ASTM D257)

Table 4 shows the electrical insulating properties of BeO heat-dissipating sheets at various BeO contents. At 21.4% BeO contents, the heat-dissipating sheets had the maximum resistance ( $2.6 \times 10^{12} \Omega \square^{-1}$ ). However, with 23.4% BeO content, BeO heat-dissipating sheets have the minimum electrical resistivity ( $1.5 \times 10^{12} \Omega \square^{-1}$ ). The electrical resistivity measurements are all within the normal range. Both PU and BeO have a high electrical resistance could be the reason. As a result, mixed PU and BeO had a higher resistance.<sup>50–52</sup> So, it concluded that the BeO heat-dissipating sheet's electrical insulating qualities were outstanding.

## Conclusions

In summary, PVA/ $\text{BeSO}_4$ /PEI precursor NFs were successfully formed by electrospinning technique mixing  $\text{BeSO}_4 \cdot 4\text{H}_2\text{O}$  salt and PEI in PVA aqueous solution, then calcination at different temperatures to produce BeO NF mats. Furthermore, the crystallinity increased dramatically as the calcination temperatures increased from 600  $^{\circ}\text{C}$  to 1300  $^{\circ}\text{C}$ . The calcination temperatures also influenced the specific surface area and average pore area

due to the removal of organics PVA. This is beneficial to improving the low specific surface area of precursor PVA/ $\text{BeSO}_4$ /PEI NFs of  $36.3 \text{ m}^2 \text{ g}^{-1}$  to  $145 \text{ m}^2 \text{ g}^{-1}$  for BeO NFs calcined at 600  $^{\circ}\text{C}$ . On the contrary, a very high calcination temperature of 1300  $^{\circ}\text{C}$  will decrease specific surface area ( $5.1 \text{ m}^2 \text{ g}^{-1}$ ) due to sintering by metal particles. This study also investigated the thermal conductivity of PU/BeO NFs composite sheets by adding PU with BeO NF mats. The heat-dissipating sheets showed improved thermal conductivity ( $1.4\text{--}14.4 \text{ W m}^{-1} \text{ K}^{-1}$ ) at 16.6–80.4 vol% BeO NFs content. The possible reason is that the elongated shape of BeO NFs serves as an effective filler, creating linear heat pathways in the resin. In addition, high-performance heat-dissipating sheets showed excellent electrical insulating properties ( $1.6 \times 10^{12} \Omega \square^{-1}$ ). This processing technique is very effective in synthesizing BeO NFs for the first time. Electrically insulated thermal conductive PU/BeO NF composite sheets would effectively dissipate heat from electronic gadgets.

## Author contributions

Md. Shakhawat Hossain: methodology, writing – original draft, investigation, data curation, visualization. Anamul Hoque Bhuiyan: experimental implementation and apparatus use. Koji Nakane (corresponding author): conceptualization, resources, design the research, formal analysis, editing, supervision.

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## Conflicts of interest

There are no conflicts of interest declared by the authors.

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## References

- 1 H. Chen, V. V. Ginzburg, J. Yang, Y. Yang, W. Liu, Y. Huang, L. Du and B. Chen, *Prog. Polym. Sci.*, 2016, **59**, 41–85.
- 2 F. Yuan, W. Jiao, F. Yang, W. Liu, Z. Xu and R. Wang, *RSC Adv.*, 2017, **7**, 43380–43389.
- 3 X. Pu, H.-B. Zhang, X. Li, C. Gui and Z.-Z. Yu, *RSC Adv.*, 2014, **4**, 15297–15303.
- 4 F. Nazeer, Z. Ma, Y. Xie, L. Gao, A. Malik, M. A. Khan, F. Wang and H. Li, *RSC Adv.*, 2019, **9**, 17967–17974.
- 5 H. Shen, J. Guo, H. Wang, N. Zhao and J. Xu, *ACS Appl. Mater. Interfaces*, 2015, **7**, 5701–5708.



- 6 Y. Xu, D. D. L. Chung and C. Mroz, *Composites, Part A*, 2001, **32**, 1749–1757.
- 7 C. Huang, X. Qian and R. Yang, *Mater. Sci. Eng. R Rep.*, 2018, **132**, 1–22.
- 8 A. R. J. Hussain, A. A. Alahyari, S. A. Eastman, C. Thibaud-Erkey, S. Johnston and M. J. Sobkiewicz, *Appl. Therm. Eng.*, 2017, **113**, 1118–1127.
- 9 X. Chen, Y. Su, D. Aydin, D. Reay, R. Law and S. Riffat, *Energy Build.*, 2016, **125**, 99–108.
- 10 W. Zhou, *Thermochim. Acta*, 2011, **512**, 183–188.
- 11 X. Lu and G. Xu, *J. Appl. Polym. Sci.*, 1997, **65**, 2733–2738.
- 12 S. Zhang, X. Y. Cao, Y. M. Ma, Y. C. Ke, J. K. Zhang and F. S. Wang, *Exp. Polym. Lett.*, 2011, **5**, 581–590.
- 13 G.-W. Lee, M. Park, J. Kim, J. I. Lee and H. G. Yoon, *Composites, Part A*, 2006, **37**, 727–734.
- 14 W. Cui, Y. Zhu, X. Yuan, K. Chen and F. Kang, *J. Alloys Compd.*, 2012, **540**, 165–169.
- 15 T. Kusunose, T. Yagi, S. H. Firoz and T. Sekino, *J. Mater. Chem. A*, 2013, **1**, 3440–3445.
- 16 G. P. Akishin, S. K. Turnaev, V. Ya. Vaispapur, M. A. Gorbunova, Yu. N. Makurin, V. S. Kiiko and A. L. Ivanovskii, *Refract. Ind. Ceram.*, 2009, **50**, 465–468.
- 17 H. Liu and S. Wu, *RSC Adv.*, 2022, **12**, 12647–12654.
- 18 M. Harada, N. Hamaura, M. Ochi and Y. Agari, *Composites, Part B*, 2013, **55**, 306–313.
- 19 K. Sato, H. Horibe, T. Shirai, Y. Hotta, H. Nakano, H. Nagai, K. Mitsuishi and K. Watari, *J. Mater. Chem.*, 2010, **20**, 2749–2752.
- 20 W. Zhou, S. Qi, Q. An, H. Zhao and N. Liu, *Mater. Res. Bull.*, 2007, **42**, 1863–1873.
- 21 S. Kume, I. Yamada, K. Watari, I. Harada and K. Mitsuishi, *J. Am. Ceram. Soc.*, 2009, **92**, S153–S156.
- 22 Y. Zhou, Y. Yao, C.-Y. Chen, K. Moon, H. Wang and C. Wong, *Sci. Rep.*, 2014, **4**, 1–6.
- 23 R. F. Hill and P. H. Supancic, *J. Am. Ceram. Soc.*, 2004, **87**, 1831–1835.
- 24 T. Zhou, X. Wang, G. U. Mingyuan and X. Liu, *Polymer*, 2008, **49**, 4666–4672.
- 25 G. P. Akishin, S. K. Turnaev, V. Y. Vaispapur, M. A. Gorbunova, Y. N. Makurin, V. S. Kiiko and A. L. Ivanovskii, *Refract. Ind. Ceram.*, 2009, **50**, 465–468.
- 26 X. Wang, R. Wang, C. Peng, T. Li and L. I. U. Bing, *Prog. Nat. Sci.: Mater. Int.*, 2010, **20**, 81–86.
- 27 S. Rezabeyk and M. Manoochehri, *RSC Adv.*, 2020, **10**, 36897–36905.
- 28 C. J. Buchko, L. C. Chen, Y. Shen and D. C. Martin, *Polymer*, 1999, **40**, 7397–7407.
- 29 J. Doshi and D. H. Reneker, *J. Electrostat.*, 1995, **35**, 151–160.
- 30 A. Ohgoshi, K. Takahashi and K. Nakane, *J. Mater. Sci.: Mater. Electron.*, 2019, **30**, 20566–20573.
- 31 K. Nakane, S. Ichikawa, S. Gao, M. Seto, S. Irie, S. Yonezawa and N. Ogata, *Sen'i Gakkaishi*, 2015, **71**, 1–5.
- 32 K. Nakane, M. Seto, S. Irie, T. Ogihara and N. Ogata, *J. Appl. Polym. Sci.*, 2011, **121**, 1774–1779.
- 33 A. Ohgoshi, S. Gao, K. Takahashi and K. Nakane, *J. Text. Eng.*, 2019, **65**, 67–72.
- 34 C. Lu, S. W. Chiang, H. Du, J. Li, L. Gan, X. Zhang, X. Chu, Y. Yao, B. Li and F. Kang, *Polymer*, 2017, **115**, 52–59.
- 35 J. M. Deitzel, J. Kleinmeyer, D. E. A. Harris and N. B. Tan, *Polymer*, 2001, **42**, 261–272.
- 36 J. Y. Lu, C. Norman, K. A. Abboud and A. Ison, *Inorg. Chem. Commun.*, 2001, **4**, 459–461.
- 37 K. H. Lee, H. Y. Kim, Y. J. Ryu, K. W. Kim and S. W. Choi, *J. Polym. Sci., Part B: Polym. Phys.*, 2003, **41**, 1256–1262.
- 38 X. Zhang, Y. Zheng, X. Feng, X. Han, Z. Bai and Z. Zhang, *RSC Adv.*, 2015, **5**, 86102–86112.
- 39 M. S. Hossain and K. Nakane, *Results Mater.*, 2022, **13**, 100241.
- 40 Y. Hou, C. Chen, K. Liu, Y. Tu, L. Zhang and Y. Li, *RSC Adv.*, 2015, **5**, 24023–24030.
- 41 A. R. Polu and R. Kumar, *Chin. J. Polym. Sci.*, 2013, **31**, 641–648.
- 42 C. Liebenow, *Electrochim. Acta*, 1998, **43**, 1253–1256.
- 43 A. B. Shatan, K. Venclíková, B. A. Zasoňska, V. Patsula, O. Pop-Georgievski, E. Petrovský and D. Horák, *Pharm. Res.*, 2019, **36**, 1–12.
- 44 D. Perra, N. Drenchev, K. Chakarova, M. G. Cutrufello and K. Hadjiivanov, *RSC Adv.*, 2014, **4**, 56183–56187.
- 45 Y. Liang, J. Ouyang, H. Wang, W. Wang, P. Chui and K. Sun, *Appl. Surface Sci.*, 2012, **258**, 3689–3694.
- 46 F. Mohandes and M. Salavati-Niasari, *RSC Adv.*, 2014, **4**, 25993–26001.
- 47 H. Lulit, S. Natnael, M. DuRe, T. Thriveni, C. Ramakrishna and A. JiWhan, *Sustainability*, 2019, **11**, 3196–3205.
- 48 C. Sarkar and S. K. Dolui, *RSC Adv.*, 2015, **5**, 60763–60769.
- 49 N. Jain, N. Marwaha, R. Verma, B. K. Gupta and A. K. Srivastava, *RSC Adv.*, 2016, **6**, 4960–4968.
- 50 Y. Mobarak, M. Bassyouni and M. Almutawa, *Adv. Mater. Sci. Eng.*, 2013, **2013**, 1–6.
- 51 A. Ş. Demirkiran, R. Artir and E. Avci, *Ceram. Int.*, 2010, **36**, 917–921.
- 52 J.-C. Zhao, F.-P. Du, X.-P. Zhou, W. Cui, X.-M. Wang, H. Zhu, X.-L. Xie and Y.-W. Mai, *Composites, Part B*, 2011, **42**, 2111–2116.

