Polymer composites based on hexagonal boron nitride and their application in thermally conductive composites

Cuiping Yu,ab Jun Zhang,b Wei Tian,*a Xiaodong Fan,a and Yagang Yao,b*

Hexagonal boron nitride (h-BN) is also referred to as “white graphite”. Owing to its two-dimensional planar structure, its thermal conductivity along and perpendicular to a basal plane is anisotropic. However, h-BN exhibits properties that are distinct from those of graphite, such as electric insulation, superior antioxidative ability, and purely white appearance. These qualities render h-BN superior as a filler in composites that require thermal conductivity while exhibiting electric insulation. Since the thermal performance of composites is mainly affected by thermal pathways, this article begins with an overall introduction of the preparation of boron nitride nanosheets, followed by a review of the fabrication of h-BN-filled composites. Lastly, the construction of thermally conductive networks is discussed.

1. Introduction

With the miniaturization and high-degree integration of modern electric devices, high performance packaging materials are urgent in thermal management to guarantee the performance and reliability of the devices.1–3 Two-dimensional (2D) materials were recently used as a new platform for thermal management applications due to their fascinating thermal properties, such as high in-plane thermal conductivity and the relatively low out-of-plane values.4 This could be ascribed to the covalent intra-layer bonds and van der Waals layer-to-layer interactions in 2D materials. For example, since graphene possesses high in-plane thermal conductivity (~5300 W m⁻¹ K⁻¹), the thermal conductivity of the target composites increased significantly with the addition of only a small amount of graphene.5,6 Black phosphorus, as a new member of 2D materials, exhibited an anisotropy thermal conductivity (~20 and ~40 W m⁻¹ K⁻¹ for the armchair and zigzag configurations, respectively).7 However, they unavoidably make the target composites electrically conductive, which could not apply in electric equipment that need thermally conductive and electric materials were recently used as a new platform for thermal management applications due to their fascinating thermal properties, such as high in-plane thermal conductivity and the relatively low out-of-plane values.4 This could be ascribed to the covalent intra-layer bonds and van der Waals layer-to-layer interactions in 2D materials. For example, since graphene possesses high in-plane thermal conductivity (~5300 W m⁻¹ K⁻¹), the thermal conductivity of the target composites increased significantly with the addition of only a small amount of graphene.5,6 Black phosphorus, as a new member of 2D materials, exhibited an anisotropy thermal conductivity (~20 and ~40 W m⁻¹ K⁻¹ for the armchair and zigzag configurations, respectively).7 However, they unavoidably make the target composites electrically conductive, which could not apply in electric equipment that need thermally conductive and electric
Hexagonal boron nitride (h-BN), also called as white graphite, is an electrical insulator with a dielectric constant 3−4. This makes it suitable for the applications in thermal management of the electronics devices and displays that are not possible for graphene and black phosphorus. Thus, inorganic ceramic h-BN were designed to fabricate various insulating composites with superior thermal conductivity.\(^7\)–\(^\text{11}\)

Boron nitride (BN) consists of boron (B) and nitrogen (N) atoms in equal amounts and is arranged in a honeycomb configuration. The materials are mainly prepared from B, boric acid (\(\text{H}_3\text{BO}_3\)), or boron trioxide (\(\text{B}_2\text{O}_3\)). BN exhibit 3 crystalline forms: h-BN (a layered structure like graphite), cubic BN (c-BN, similar to diamond), and wurtzite BN (w-BN, analogous to lonsdaleite). Among the 3 BN phases, h-BN is the most stable under standard conditions and is a sp\(^3\)-hybridized 2D-layered insulator. Within a 2D layer, the B and N atoms are alternatively bonded together by robust B-N covalent bonds, while the 2D layers are stacked with each other via weak van der Waals forces. B-N bonds are partially ionic owing to the higher electronegativity of the N atom, which varies from the pure covalent C-C bonds in a graphitic structure. The strong ionic bond of h-BN renders h-BN more difficult to exfoliate and functionalize than graphite. The crystallographic parameters of h-BN are \(a = 0.250 \text{ nm}\) and \(c = 0.666 \text{ nm}\), and the interlayer spacing is 0.333 nm. These characteristics allow h-BN to interact with graphene, enabling potential applications in many devices.

Owing to its distinct features, h-BN exhibits various properties, such as anisotropy parallel and perpendicular to a basal plane,\(^\text{12}\)–\(^\text{14}\) constant wide band gap (\(\sim 5.97 \text{ eV}\)),\(^\text{15}\) superior antioxidative and structural stabilities (high thermal stability of up to 1000 °C in air and 1400 °C in vacuum),\(^\text{16}\) high thermal conductivity (600 W m\(^{-1}\) K\(^{-1}\) for the in-plane direction, 30 W m\(^{-1}\) K\(^{-1}\) for the out-of-plane direction),\(^\text{17}\) and superior thermal expansion coefficient (1-, 2-, and 9-layer boron nitride nanosheets (BNNSs) showed the thermal expansion coefficient of about \(-3.41 \times 10^{-2}, -3.15 \times 10^{-2}\), and \(-3.78 \times 10^{-2}\) (cm K\(^{-1}\)), respectively),\(^\text{18}\) as well as superior mechanical properties (elastic constant of 220–510 N m\(^{-1}\) and Young’s modulus –1 TPa).\(^\text{19}\) These characteristics render h-BN as an excellent filler in thermally conductive while electric insulating composites.

In this review, we first describe recent advances in the preparation of boron nitride nanosheets (BNNSs) as well as the fabrication of h-BN-based composites. The construction of thermally conductive networks is then reviewed based on their different thermally conductive pathways, followed by the summary and outlook.

2. Preparation of BNNSs

BNNSs are prepared using numerous synthesis methods, most of which are similar to well-established techniques in preparing corresponding carbon materials, with minor modifications.
Among the preparation methods reported, mechanical exfoliation, liquid exfoliation, chemical vapor deposition (CVD), and epitaxy are the most widely studied. In this section, some of these methods are briefly reviewed.

Mechanical exfoliation consists of direct mechanical exfoliation and other exfoliation methods by shear forces. Ball milling is regarded as an excellent approach that uses shear forces to exfoliate bulk h-BN material down to few-layered sheets. In one study, high-yield and high-quality BNNSs were obtained by low-energy ball milling and integration of low-energy ball milling with sonication. Yao et al. exfoliated h-BN in sodium dodecyl sulfate (SDS)–water solutions into mono- and few-layered BNNSs with a concentration of about 1.2 mg mL$^{-1}$ (Fig. 1). This dispersion exhibited excellent stability (Fig. 1c). All of the aforementioned strategies were performed in liquid media. Some solid media were also reported.

Liquid exfoliation of h-BN was successfully conducted in 1,2-dichloroethane solution of poly(m-phenyl-enevinylene-co-2,5-dioxy-p-phenylenevinylene), water, SDS and water, N,N'-dimethylformamide (DMF), sulfonic acid, and methanesulfonic acid. To understand the functions of different solvents used in h-BN exfoliation, the Hansen solubility parameter (HSP) theory was proposed by Coleman et al. The exfoliation effect of the solvents could be verified by polar, hydrogen-bonding, cohesive energy density, which should minimize the energy of exfoliation or with strong, sufficient surface tension (e.g. $\gamma \approx 40$ mJ m$^{-2}$). With this theory, isopropyl alcohol (IPA) was a good exfoliation agent with a BNNS yield of $\sim$50% (0.06 mg mL$^{-1}$ BNNS concentration). On the basis of this theory, analyzing the HSP distance and $R_a$ value of diverse solvent mixtures, 55% ethanol and water solvent were better solvents than IPA that the dispersion concentration was 0.075 $\pm$ 0.003 mg mL$^{-1}$. The effect of dispersion is presented in Fig. 2. Subsequently, ammonia solution and IPA were chosen as the exfoliation agents; ammonia molecules could intercalate the h-BN layers and thus benefited the exfoliation process. Given the large surface tension ($\gamma = 33.3$ mN m$^{-1}$) and small HSP distance ($R_a = 2.59$ A) of thionyl chloride, it could be potentially used to exfoliate and stabilize BNNS dispersions.

Epitaxial growth of monolayer h-BN was achieved on Pt (111) and Ru (001) substrates via adsorption and decomposition of borazine. Single-crystalline monolayer h-BN on the Ni (111) substrate with borazine as the precursor was prepared by Oshima et al. The Ni foil was considered as a preferred substrate because of the minor difference in the lattice constant between the Ni (111) surface (2.49 A) and bulk h-BN (2.5 A). Alternatively, B-trichloroborazine (ClBNH)$_3$ was found to be a better precursor for monolayer h-BN growth on Pt (111) and Ni (111). BNNSs in the form of a so-called nanomesh were synthesized on a Rh (111) substrate by Corso et al. The types of substrate influenced the pore shapes and sizes of the nanomesh. A nanomesh could be grown on Ru (0001) and Rh (111). However, a flat monolayer h-BN could be grown on Pt (111), Cu (111), Fe (110), Pd (111), Cr (110), Mo (110), and Ni (111). Monolayer and nanomesh h-BN coexisted when Ir (111) was used as the substrate. Monolayer h-BN was grown on a four-inch Rh (111) substrate in a modified ultrahigh vacuum chamber. This h-BN surface exhibited a corrugated morphology. Similar patterns were observed on an epitaxial h-BN layer measuring 2 in. The wrinkle-free h-BN could be grown on a c-plane sapphire substrate measuring 2 in with the AA stacking order by modified low-pressure chemical vapor deposition (CVD).

Cu and Ni foils were traditionally used as substrates for 2D h-BN growth because of their low-cost and excellent catalytic properties. Typical CVD processes for BNNS growth have used borazine and ammonia borane with Pt, Ni, and Cu as substrates. Wang et al. prepared a triangular monolayer single crystal h-BN measuring 20 $\mu$m. This size varied with the thermal annealing time of Cu foils (Fig. 3). Several catalyst-based substrates, such as heteroepitaxial Cu–Ni alloy, Co film, and Si-doped Fe substrate, have been explored. A study found that nanocrystalline graphene could be used as a seed layer for synthesizing large-area h-BN. In addition, h-BN could be grown on insulating substrates, such as SiO$_2$/Si without a catalyst, via microwave plasma CVD. Few-layered BNNSs were grown by using a mixture of B, magnesium oxide (MgO), ferrous oxide (FeO) powders, and ammonia on silica (SiO$_2$)/Si. Wafer-scale multilayer BNNSs were synthesized in vacuum with a subnanometer scale surface roughness.

3. Fabrication of h-BN-filled composites

The processing step exerts significant effects on the bulk properties of composites. Thus, methods of fabricating h-BN-filled composites are briefly reviewed, including solution mixing, melt blending, in situ polymerization, and ball milling.
3.1 Solution mixing

Solution mixing is a facile-operated, commonly used technique for fabricating h-BN-filled polymer composites. A suitable solvent is crucial for the uniform dispersion of h-BN and the polymer matrix, forming metastable suspensions of h-BN or h-BN/polymer mixtures. The critical technique during this process involves uniform stirring, which often requires the use of high-power ultrasonication. Some functionalized strategies, such as heat treatment, acid treatment or h-BN functionalization, were conducted to improve the dispersion of h-BN. Many h-BN/polymer composites were successfully prepared by solution mixing, including BN/polymethyl methacrylate (PMMA), BN/poly (vinyl alcohol) (PVA), BN/polyethylene terephthalate (PBT), BN/epoxy, BN/polyurethane acrylate (PUA), and BN/silicone rubber (SiR). Song et al. fabricated BNNS/PVA films by stirring the mixture of aqueous PVA solution and IPA suspension of BNNSs. The solvent was removed with vigorous stirring and then subjected to casting on an etched glass slide, resulting in a thin film with a thermal conductivity higher than 30 W m⁻¹ K⁻¹ (Fig. 4). However, solution mixing is limited to polymers that can be dissolved in solvents which are also suitable for the preparation of a stable suspension of h-BN.

Fig. 2 (a) Photographs of BN dispersions in several ethanol/water mixtures stored under ambient conditions for 1 week. (b) BN suspensions in various ethanol/water mixtures and calculated $R_g$ values are denoted by dots and solid lines, respectively. Reproduced with permission from ref. 35.

Fig. 3 (a) Dissociation of ammonia borate and h-BN growth. SEM images of h-BN grown without pre-annealing (b) and with annealing for 3 h (c) and 6 h (d). (e) Photograph showing variations in the domain size distributions of h-BN with the annealing time of Cu foils. SEM images of h-BN film formed by h-BN domains with growth times of about 65 min (f) and 70 min (g). Reproduced with permission from ref. 61.
3.2 Melt blending

Melt blending is a solvent-free process suitable for industrial processes. This technique is usually applied to disperse sheets in a thermostatic polymer matrix by using high-temperature and high-shear forces. h-BN/bisphenol E cyanate ester (BECy) nanocomposites were fabricated by preheating dopamine-treated h-BN nanoparticles at 130 °C and then mixing with a BECy resin, followed by curing at 180 °C and 250 °C (Fig. 5). However, melt blending was generally less effective in dispersing sheets in polymers, compared with solution mixing. This approach was limited to low filler loadings because of the high viscosity of the composites at high sheet loadings. Xie et al. subsequently prepared the mixture of BNNSs with poly(ethylene terephthalate) by melt processing. However, the BNNSs preferred to restack during the drying process. Thus, Guo et al. developed a slurry compounding strategy to fabricate BNNS/styrene–butadiene rubber (SBR) composites by using a conventional two-roll mill. The BNNS slurry was prepared using a solvent exchange route. Silane was added to strengthen the interfaces of BNNSs and SBR matrix. Compared with the neat SBR composites, the composites for 10.5 vol% Si–BNNS loading increased the thermal conductivity by 253%.

3.3 In situ polymerization

Usually, a high filler loading would sacrifice the mechanical property of the composites. This could be ascribed to the poor stress transfer from matrix to filler caused by agglomeration of fillers. Thus, in situ polymerization was conducted to increase the interface bonding or adhesion between the fillers and polymer matrix. In situ polymerization involves dispersing the reactive monomer into sheet layers, followed by polymerization. This technique allows covalent bonding between functionalized sheets and the polymer matrix via chemical reaction. Distinct composites were prepared using this technique. Styrene was initiated on modified BNNSs by reversible addition–fragmentation chain transfer polymerization to prepare BNNS/polystyrene (PS) composites. Composites of h-BN/PMMA were fabricated by dispersing h-BN particles into methyl methacrylate monomers by bulk polymerization and atom transfer radical polymerization. A typical example was
reported by Lee et al.\textsuperscript{89} in which poly(caprolactone) (PCL) was chemically grafted onto BNNSs by oxidized BNNSs (oxi-BNNSs) and subsequent ring-opening polymerization of PCL (Fig. 6). The resulting composites exhibited a thermal conductivity of 1.96 W m\textsuperscript{-1} K\textsuperscript{-1} with the incorporation of 20 wt% BNNSs.

3.4 Ball milling

Ball milling is a method of grinding materials into extremely fine powders or dispersions for practical application. The milling balls used are made of different materials, including stainless steel, ceramic, flint pebbles, and so on. High pressure is generated during milling because of the collision of tiny rigid balls in a concealed container. The behavior of powder particles upon milling is generally distinguished into mechanical milling, mechanical alloying, and mechanochemical synthesis. High-energy ball milling presents inherent advantages, such as low cost, versatility, and scalability. Therefore, high-energy ball milling is considered an industrialized technique for manufacturing materials in large quantities. Ball milling on layered materials generates strong shear and compression forces, which can peel off and cleave the materials (Fig. 1).\textsuperscript{19} Thus, ball milling is an excellent approach to reducing the thickness of h-BN. Lee et al.\textsuperscript{39} recently fabricated high-quality and high yield hydroxide BNNSs (OH–BNNSs) with high efficiency by hydroxide-assisted ball milling. The resulting OH–BNNSs showed little damage, relatively large sheets with an average size of 1.5 \( \mu \)m, and a high yield of 18%. Functionalized BNNSs (f-BNNSs) were used to fabricate polyethylene composites. Oxygen permeability was reduced by 46%, and water vapor permeability decreased by 34% with 5 wt% f-BNNS loading (Fig. 7). Various parameters, such as ball-to-powder ratios, milling speed, milling ball sizes, and milling agents, considered in the preparation of BNNSs in benzyl benzoate were systematically investigated by Chen et al.\textsuperscript{24} The obtained sample had a diameter ranging from 0.5 \( \mu \)m to 1.5 \( \mu \)m and a yield of 13.8%.

4. Conducting thermally conductive networks

The effective routes to assemble fillers in the polymer in order to construct well-organized and interconnected thermally conductive networks are critical for heat transfer. The usual strategies conducted for the construction of thermally conductive networks exhibit alignment, a three-dimensional (3D) network, an interface effect, a hybrid effect, and so on.

4.1 Alignment

h-BN is a platelet-shaped particle with a high aspect ratio. In one study, the thermal conductivity of h-BN reached 600 W m\textsuperscript{-1} K\textsuperscript{-1} in the in-plane direction and 30 W m\textsuperscript{-1} K\textsuperscript{-1} in the out-of-plane direction.\textsuperscript{22} Owing to the anisotropic shape and thermal conductivity of h-BN, the arrangement of h-BN in the polymer matrix considerably affected the thermal property of the h-BN-filled composites. Numerous reports recently evaluated the influence of orientation on thermal conductivity enhancement in a preferential direction. The methods used to orient h-BN included processing and external forces.

4.1.1 Alignment during processing. As shear force is introduced by processing methods, such as oscillatory shearing, tape casting, and injection molding, h-BN typically tends to align parallel to the planer direction, as in the case of h-BN oriented in PVA composite films by tape-casting.\textsuperscript{74}
thermal conductivity reached 5.4 W m\(^{-1}\) K\(^{-1}\) for 10 vol% filler contents. Ceramic particles were oriented in polyvinyl butyral and PVA matrices prepared using a similar method.\(^\text{92,93}\) Kuang et al.\(^\text{14}\) fabricated BNNS/SiR and BNNS/natural rubber (NR) composites with ultrahigh degree of orientation because of strong shear forces introduced by a two-roll mill (Fig. 8). The thermal conductivity of the composites was ~4 W m\(^{-1}\) K\(^{-1}\) with 23 vol% BNNSs in SiR composites. h-BN/Polypropylene
composites with a high degree of orientation were obtained by injection molding. The corresponding thermal conductivity showed a large increase, which reached 2.1 W m$^{-1}$ K$^{-1}$ with the incorporation of 40 vol% h-BN.

4.1.2 Alignment by external fields. Alignment of h-BN in composites can be achieved by applying a suitable external field, such as magnetic, electric, and gravitational force fields.

4.1.2.1 Magnetic field alignment. Studart et al. proposed in 2012 an approach to align anisotropic reinforcing microparticles in the composite matrix by applying external magnetic fields. Reinforcing particles are often diamagnetic; as such, they were first coated with superparamagnetic nanoparticles. The filler arrangement in low-viscosity suspending fluids was controlled using a low external magnetic field. h-BN complex sheets modified with iron oxide nanoparticles were successfully reoriented in the direction vertical to the film plane in the study by Cho et al., as shown in Fig. 9. Thus, iron oxide-coated h-BN filled epoxy composites exhibited excellent thermal properties. The highest thermal conductivity obtained was 4.7 W m$^{-1}$ K$^{-1}$ at 20 vol% iron oxide-coated h-BN for an epoxy resin system. Luo et al. used a similar method to fabricate iron oxide-coated h-BN/SiR composites. The thermal conductivity of the composites was 44.5% higher than that of the unaligned composites with 9.14 vol% filler loading. A theoretical model was used to analyze the results, suggesting that the orientation of the BNNSs mainly led to the improvement of thermal conductivity. CVD was also conducted to deposit iron oxide onto BNNSs. The obtained modified BNNSs were successfully oriented in the polysiloxane matrix with a magnetic field (Fig. 9).

4.1.2.2 Electric field alignment. BNNSs can be oriented by an extra electric field. BNNS/polysiloxane composites were fabricated under a dc electric field (Fig. 10). BNNSs in the matrix are reoriented in the electric field direction. This direction can be controlled to be perpendicular to the film plane. Field-induced orientation of BNNSs in the polysiloxane matrix was successfully accomplished.

4.1.2.3 Gravitational force-induced alignment. Gravitational force is preferred to align BNNSs in the in-plane direction. This strategy was facile, simple, and resulted in a high degree of orientation. The highest thermal conductivity reported in h-BN-filled composites was 145.7 W m$^{-1}$ K$^{-1}$ at 50 wt% h-BN content, which was ascribed to the constitution of hierarchically
arranged 2D BNNSs and 1-dimensional (1D) cellulose formed during vacuum-assisted filtration. A large contact area was formed between BNNSs in this structure, thereby minimizing the influence of interface thermal resistance along the thermally conductive direction. An orderly oriented structure based on cellulose nanofibers (CNFs) and f-BNNSs was prepared by simple vacuum-assisted filtration with superior thermal conductivity of up to 30.25 W m\(^{-1}\) K\(^{-1}\) with the incorporation of 70 wt% f-BNNS (Fig. 11). In addition, pure laminates of h-BN could be prepared by vacuum filtration of the h-BN dispersion prepared via liquid exfoliation in IPA. The thermal conductivity of such insulating laminates reached 20 W m\(^{-1}\) K\(^{-1}\). By improving the quality of the flake-to-flake interface, the interface thermal resistance decreased and thus enhanced thermal conductivity. OH–BNNSs could be obtained by molten alkali-assisted exfoliation of h-BN, which improved the interfacial bonding strength between BNNS layers. Subsequent filtration resulted in OH–BNNS papers with superior thermal conductivity of 58 W m\(^{-1}\) K\(^{-1}\). The traditional platelet filler contents in the polymer composites were lower than 40 wt%. However, a high filler loading can be achieved by vacuum-assisted filtration. Composite paper analogous to artificial nacre consisting of noncovalent functionalized BNNSs (NF-BNNSs) and PVA with ultrahigh filler loading of 94 wt% was fabricated by vacuum-

---

**Fig. 8** (a) Preparation of BNNS/SiR and BNNS/NR composites. High-resolution TEM images of BNNS/SiR (b) and BNNS/NR (c) composites with 4 vol% BNNS loadings. Thermal conductivities of BNNS/SiR (d) and BNNS/NR (e) composites with different BNNS contents. Reproduced with permission from ref. 14.

---

RSC Advances

Review

RSC Adv., 2018, 8, 21948–21967

This journal is © The Royal Society of Chemistry 2018

Open Access Article. Published on 14 June 2018. Downloaded on 11/19/2023 12:29:18 PM.

This article is licensed under a Creative Commons Attribution-NonCommercial 3.0 Unported Licence.
induced filtration.\textsuperscript{78} A framework constructed by hierarchically arranged BNNSs and PVA was formed by gravitational force. This paper exhibited high thermal conductivity of 6.9 W m\(^{-1}\) K\(^{-1}\).

**4.2 3D networks**

Isotropic thermally conductive bulk composites are favorable to a certain extent in some applications. Thus, forming three-dimensional (3D) networks with high quality and low thermal resistance for heat conduction present a significant challenge. Up to now, the commonly strategy used to construct 3D networks was to form thermally conductive networks firstly and then infiltrating with polymer. Thus, the thermal performance of the composites improved a lot at only a low filler loading. The thermal conductivity enhancement efficiency of 3D network composites to pure polymer matrix was expressed by thermal conductivity enhancement per 1 vol% loading \((\eta)\), which was defined as \(\eta = (K - K_m) \times 100\% / (100 \times VK_m)\), where \(K\) and \(K_m\) represented the thermal conductivity of the composites and polymer matrix, respectively. And \(V\) represented the volume contents of h-BN in composites. As shown in Fig. 12, CNF-supported 3D-interconnected BNNS (3D-C-BNNS) aerogels were fabricated by self-assembly of BNNSs on a 3D cellulose skeleton via sol–gel and freeze-drying.\textsuperscript{106} Composites exhibiting high thermal conductivity were then prepared by impregnating epoxy resin into the 3D BNNS aerogel. Ultrahigh thermal conductivity of 3.13 W m\(^{-1}\) K\(^{-1}\) was achieved at a low BNNS loading level of 9.6 vol\%, which corresponded to a thermal conductivity enhancement of about 1400\%. In addition, the thermal conductivity enhancement could be easily adjusted by tailoring the BNNS loadings in the interconnected 3D network.

An ice-templated approach was reported to construct 3D BNNS networks (Fig. 13).\textsuperscript{105} This strategy obtained epoxy composites with high thermal conductivity (2.85 W m\(^{-1}\) K\(^{-1}\)) at a relatively low filler content (9.29 vol\%). Yang et al.\textsuperscript{107} subsequently used a similar method to fabricate graphene oxide (GO)/h-BN hybrid porous scaffolds by introducing polyethylene glycol and obtained thermal conductivity of 1.84 W m\(^{-1}\) K\(^{-1}\) at 19.2 wt% h-BN.

**4.3 Interface effect**

A large body of evidence supports the finding that surface modification can reduce interface thermal resistance. For chemical surface modification, covalent bonding or
Fig. 10  (a) Illustration of the experimental setup for the fabrication of aligned BN/polymer films. Cross-sectional SEM micrographs of an oriented BN/polysiloxane composite at 10 vol% BN content without (b) and with (c) the application of electric fields. Reproduced with permission from ref. 99.

Fig. 11  (a) Demonstration of BNNSs or f-BNNSs in CNF aqueous solution during vacuum-assisted filtration. (b) Thermal conductivity of CNF composite films. (c) Photographs of CNF/BNNS and CNF/f-BNNS films with 70 wt% BNNS loadings and the corresponding SEM images. (d) In-plane thermal conductivity of CNF/f-BNNS films at 70 wt% filler contents with different thicknesses. Reproduced with permission from ref. 102.
to have improved the compatibility among fillers and the matrix, resulting in a high thermal conductivity of 3.92 W m\(^{-1}\) K\(^{-1}\) at 10 wt% h-BN loadings in the PVA matrix. Tang et al.\(^\text{109}\) employed a mixed solution of concentrated sulfuric acid (H\(_2\)SO\(_4\)) and nitric acid (HNO\(_3\)) to synthesize acid-treated BNNSs to enhance the thermal performance of epoxy composites. Thus, the thermal conductivity channel of the obtained composites was first constructed at 5.05 wt%, which was higher than that of pure BNNS-epoxy composites at 6.10 wt%. These methods were based on a liquid medium, which was difficult to remove completely because of their high boiling points. Therefore, a 1-step route in the preparation of OH–BNNS was developed by Huang et al.\(^\text{77}\) by which h-BN could be directly exfoliated at a high temperature of 850 °C (Fig. 14). The resulting OH–BNNSs exhibited excellent dispersibility in water (concentration of 0.3 mg mL\(^{-1}\)) and ethanol (concentration of 0.06 mg mL\(^{-1}\)), rendering the OH–BNNSs effective as a hydrogel additive. The OH–BNNS/poly(N-isopropylacrylamide) (PNIPAM) hydrogel composite showed a 41% increase in thermal conductivity at 0.07 wt% OH–BNNS. In addition, Xu et al.\(^\text{75}\) treated h-BN with acetone, HNO\(_3\), H\(_2\)SO\(_4\), and silane, showing varying levels of effectiveness in enhancing thermal conductivity. The results exhibited that surface treatment using silane Z-6040 (2.4%) was most effective for h-BN. The thermal conductivity reached 10.3 W m\(^{-1}\) K\(^{-1}\) at 57 vol% h-BN loading. The introduced –OH groups improved the interface compatibility; thus, the diffuse path of phonon was opened between BNNSs and the polymer matrix, leading to increased thermal conductivity.

The OH–BN or NH\(_2\)–BN typically served as the initiators in the design of many complex h-BN derivatives. The obtained h-BN derivatives were used as fillers to enhance the interface adhesion between h-BN and the polymer matrix, leading to a higher dispersion of platelets in polymers and enhancing the thermal performance. To enhance the affinity of the BNNS/polymer interface, a hyperbranched polyamide (HBP) was attached to the modified BNNSs. The HBP was formed by introducing amino groups to the surface of BNNSs and then initiating HBP growth (Fig. 15).\(^\text{75,84}\) The obtained epoxy composites exhibited a thermal conductivity of 9.81 W m\(^{-1}\) K\(^{-1}\) with 50 vol% BNNSs. Kim et al.\(^\text{81,84,118}\) conducted studies on grafting silane coupling agents onto the h-BN surface for a high thermal performance. The common strategy was to attach a hydroxyl group onto the h-BN surface by using sodium hydroxide at a high temperature, followed by grafting silane coupling agents, such as 3-(glycidoxypropyl)trimethoxysilane, 3-chloropropyltrimethoxysilane,\(^\text{81,110}\) and 3-aminopropyltriethoxysilane,\(^\text{81}\) to modify the surfaces, which was a smart route to reduce interface thermal resistance. The chemical reagents used in these strategies were not environmentally friendly, and solvent removal entailed difficulty. Consequently, a solvent-free method was developed by Zhi et al.\(^\text{111}\) to fabricate epoxy composites. The h-BN powder was treated with KH550, a silane coupling agent, followed by mixing with a solid-state epoxy resin. This strategy was suitable for nearly any filler fraction. The maximum thermal conductivity reached 5.24 W m\(^{-1}\) K\(^{-1}\).

**4.3.1 Covalent modification.** Covalent functionalization of BNNSs requires the formation of covalent bonds on the basal plane of the BNNS lattice. Attributed to the partial ionic characteristic of B–N bonds in an h-BN structure, its B atom bore a partially positive charge, whereas the N atom bore a negative charge. This characteristic exposed B and N sites to attacks by nucleophilic and electrophilic groups, respectively.

Several studies have thus far been reported on the production of OH–BNNSs and subsequent OH–BNNSs/polymer composites with enhanced thermal performance. In 2013, 30% hydrogen peroxide was used to treat h-BN powders in an autoclave at 100 °C.\(^\text{108}\) The functionalization of h-BN was found noncovalent bonding are the commonly used methods to connect the polymer matrix and fillers.
4.3.2 Noncovalent modification. Noncovalent bonding is a facile method to modify the surface without introducing defects on the surface of BNNSs. Noncovalent modification was the technique employed to physically adsorb polymer on the surface of BNNSs. Thus, the crystallinity of noncovalent functionalized BNNSs was perfectly maintained even after treatment while interfacial adhesion between BNNSs and the matrix increased. Therefore, the thermal conductivity of the composites was enhanced.

Owing to its superior adhesive ability, polydopamine (PDA) was typically applied to modify the surface of h-BN. Xu et al.\textsuperscript{74} adopted dopamine chemistry to coat a thickness of about 4 nm PDA on the h-BN surface (Fig. 16). The thermal conductivity of h-BN@PDA/PVA composites was about 5.4 W m\textsuperscript{-1} K\textsuperscript{-1} with the addition of 10 vol\% h-BN@PDA. Meanwhile, Kessler et al.\textsuperscript{73} used a similar method to prepare h-BN@PDA/BECy composites and achieved a lower thermal performance of 0.5 W m\textsuperscript{-1} K\textsuperscript{-1} for a 15 vol\% filler loading. In addition, chlorosulfonic acid (CSA) exhibited strong physical adsorption on h-BN surfaces.\textsuperscript{80} The physical adsorption of CSA provided high solubility for BNNSs in various organic solvents and thermoplastic polymers, yielding NF-BNNS/PBT composite films with enhanced through-plane thermal conductivity of up to 11.0 W m\textsuperscript{-1} K\textsuperscript{-1}.

4.4 Hybrid effect

The simultaneous use of fillers with different shapes and types was an alternative strategy to enhance the thermal performance of composites.

4.4.1 Morphology and size effect. The effects of 11 artificially designed filler shapes on the thermal conductivity of the composite materials were evaluated.\textsuperscript{112} The results indicated that the best fillers have long heat transfer distances and continuously large contact areas. Double Y-shaped fillers were considered preferable to other fillers, whereas spherical fillers were considered the worst. In addition to shape, hybrid sizes can influence thermal performance because large h-BN has a longer heat transfer distance than that of small h-BN. The large h-BN contributed more to the thermal conductivity of the material. The small h-BN could bridge the distances between h-BN crystals to increase contact.\textsuperscript{113,114} Hybrid h-BN with M\textsubscript{0.3} μm : M\textsubscript{6} μm : M\textsubscript{20} μm at a mass ratio of 1 : 1 : 2 in a total of 40 vol\% leads to a higher thermal conductivity of 1.482 W m\textsuperscript{-1} K\textsuperscript{-1} in SiR composites than that of 50 vol\% BN\textsubscript{6} μm with a thermal conductivity of 1 W m\textsuperscript{-1} K\textsuperscript{-1}.\textsuperscript{115} Subsequently, microsized and nanosized BN were combined to prepare polyimide (PI) films.\textsuperscript{116} The obtained sample filled with 30 wt\% h-BN (microsize : nanosized = 7 : 3 wt\%) obtained a thermal conductivity of up to 1.2 W m\textsuperscript{-1} K\textsuperscript{-1}. This value was higher than that obtained using pure microsized BN. These enhancements in thermal property were attributed to the high
packing density and ease of formation of heat transfer pathways. The studies mainly discussed the effects exerted by microsized BN, whereas those by nanosized BN were rarely reported. Thus, Kemaloglu et al. used 3 micro-size and 2 nanosized h-BN with different particle sizes and shapes in SiR composites. The results showed that nanosized BN could not promote the thermal conductivity as much as the microsize BN. A plate-like particle with an aspect ratio of 20 was believed to exhibit the highest thermal performance.

Fig. 14 (a) Exfoliation and hydroxylation of h-BN. (b) TEM image showing a Moiré patterns. (c) AFM observation showing OH–BNNSs and corresponding height profiles. (d) TEM image showing few-layered OH–BNNSs and the corresponding SAED. (e) Heat transfer in OH–BNNS/PNIPAM hydrogel. (f) Thermal conductivity variations of hydrogels with OH–BNNS (inset: percentages of thermal enhancement). Reproduced with permission from ref. 71.
4.4.2 h-BN/nanoparticle-filled composites. Interfacial thermal resistance is regarded as the main factor influencing the thermal performance of composites, which arises from poor mechanical or chemical adherence at interfaces. The aforementioned strategies were related to the reduction in filler/matrix resistance. However, studies on the filler/filler interface have rarely been reported. Jiang et al.\textsuperscript{118} used $\alpha$-alumina ($\alpha$-Al$_2$O$_3$) to bridge BNNSs to construct more compactible 3D thermally conductive networks, resulting in more heat transfer pathways in the composites. The obtained sample exhibited a thermal conductivity of 0.808 W m$^{-1}$ K$^{-1}$ at 26.5 vol% filler loading. Permal et al.\textsuperscript{119} recently combined polygonal Al$_2$O$_3$ and BN to improve the thermal performance of epoxy composites, which reached 0.57 W m$^{-1}$ K$^{-1}$ with the addition of a 30 wt% filler (Al$_2$O$_3$ : BN$_{\text{filler}}$ = 5 : 5). Layer-shaped fillers were more favorable than the ball-shaped or sharp-corner-shaped fillers in improving the thermal performance of epoxy resins.\textsuperscript{120} However, the thermal conductivity remained low because of the absence of contact between Al$_2$O$_3$ and BNNSs. Silver nanoparticles (AgNPs) could be sintered together during epoxy curing and acted as “solders” to link BNNSs. AgNP-deposited BNNSs (BNNS/AgNPs) were used as fillers for the epoxy matrix and provided a higher thermal conductivity of 3.06 W m$^{-1}$ K$^{-1}$ with 25.1 vol% BNNS loading.\textsuperscript{121} However, the obtained thermal conductivity remained lower than 5 W m$^{-1}$ K$^{-1}$. Thus, silicon carbide nanowires (SiCNWs) were introduced to enhance the thermal performance of PVA composites.\textsuperscript{122} Bio-inspired BNNS–AgNP/SiCNW–AgNP/PVA composite papers were fabricated and exhibited a superior thermal conductivity of about 21.7 W m$^{-1}$ K$^{-1}$. The preparation process is illustrated in Fig. 17.
4.4.3 h-BN/carbon material-filled composite. To enhance the affinity between fillers and the polymer matrix, the common route was to modify the surface of the fillers. Carbon materials, such as graphene and carbon nanotubes (CNTs), and h-BN were combined as a mixed filler to improve the thermal performance of the polymer matrix. The mixed fillers were found to exert a synergistic effect on the thermal enhancement of the composites. When coupling agent-functionalized BN (f-BN) was combined with glycidyl methacrylate-grafted graphene (g-TrG) and then added into the PI matrix, the voids between BN and the PI matrix were filled by the graphene nanolayers. The process is illustrated in Fig. 18. The resulting composites containing additional 1 wt% g-TrG exhibited a thermal conductivity of 2.11 W m⁻¹ K⁻¹, which was at least doubled to the value of f-BN/PI. The graphene content in the mixed fillers was smaller. Meanwhile, when the BN content was smaller, the nanosized BN was filled into the graphene interspaces and thus formed a BN/graphene stacked structure, markedly enhancing the thermal conductivity. Additional 1.5 wt% h-BN could increase the thermal conductivity to 38% and 34% in PS and polyamide 6 (PA 6) matrix, respectively. The previously described fillers were based on 2D platelets. Once the 1D and 2D fillers were mixed, a hybrid 3D filler network structure could be formed. Thus, a small amount of CNTs (2 wt%) was introduced into the BN-filled poly(vinylidene fluoride) (PVDF) composites. The results showed that the addition of CNTs to the BN/PVDF composites facilitated the formation of a denser 3D BN/CNT network structure. The thermal conductivity of 1.3 W m⁻¹ K⁻¹ at 22 wt% filler loading (2 wt% CNTs and 20 wt% BN) was thus enhanced. This thermal conductivity was higher than that of PVDF/20 wt% BN (0.97 W m⁻¹ K⁻¹) and PVDF/2 wt% CNTs (0.28 W m⁻¹ K⁻¹).

Fig. 16 (a) Process of dopamine chemistry. (b) TEM image of h-BN@PDA microplatelets. (c) Fracture SEM image of the h-BN@PDA composite film with 20 vol% h-BN contents prepared by doctor blading. (d) Thermal conductivity of the fabricated composite films. Reproduced with permission from ref. 74.

Fig. 17 (a) Fabrication of BNNS–Ag/SiCNW–Ag/PVA paper. TEM images of BNNS–Ag (b) and SiCNW–Ag (c). (d) Fracture SEM observation of BNNS–Ag/SiCNW–Ag/PVA paper. (e) Thermal conductivity of BNNS/PVA and BNNS–Ag/SiCNW–Ag/PVA papers. Reproduced with permission from ref. 122.
provide a high quality. However, the high temperature usually smaller than other methods, such as CVD. Thus, it was desirable to develop an effective method to promote the dispersion of fillers in the matrix as well as maintain the quality of the fillers. Third, the strategies for constructing thermally conductive pathways have rarely been reported. Owing to the anisotropic thermal performance of 2D h-BN, the orientation preferred to prepare a composite with a high thermal conductivity in a special direction. 3D networks in perpendicular and parallel directions could lead to a high thermal conductivity. However, the strategies for constructing orientation or 3D networks remained limited. In addition, mixing hybrid fillers was a significant method used to form and improve heat conduction. However, the interface between nanoparticles and BNNS has great influence on the heat transfer in the hybrids. Therefore, the synergistic effect between BNNS and other functional materials, especially different polymers and various nanoparticles, remained to be exploited in the near future.

5. Summary and outlook

In the current study, the fabrication methods of BNNSs and h-BN-filled composites, as well as the construction of thermally conductive networks, were summarized based on the most updated research. Studies on BNNS-filled thermally conductive polymer composites remained at their early stage for several reasons. First, the preparation of BNNSs on a large scale was difficult for a strong “lip–lip” interaction between B–N bonds on the BNNS plane that was difficult to exfoliate. Several techniques, such as mechanical and liquid exfoliation, CVD, and epitaxy, were conducted to fabricate 2D h-BN with configurations and variations in quality. Liquid exfoliation by soninating bulk h-BN in suitable solvents would lead to a high yield. However, the lateral size of BNNS obtained by this method was usually smaller than other methods, such as CVD. Thus, it was desirable to develop effective strategies to prepare BNNS with large lateral size. The sample obtained by CVD growth could provide a high quality. However, the high temperature requirement and the transfer process, as well as the use of catalytic substrates, could be the main restrictions for the production of highly crystalline h-BN and construction of high-performance devices. Second, the properties of the interface and dispersion exerted important effects on the composite materials. The BN nanostructure was difficult to disperse uniformly in the polymer matrix due to its inert surface. Generally, the universal method used to improve the interface between BNNSs and the polymer matrix was surface modification. The common groups were identified as the hydroxyl group and the amino group. The hydroxyl group was connected to the B site, whereas the amino group was hydrogenated at the N site. Surface modification of h-BN was an optimal method to decrease the interface thermal resistance among the fillers and the matrix, as well as to promote good dispersion. In addition, since ball milling was based on mechanochemistry, it was found to be a good protocol to improve the interface and dispersion between fillers and matrix. However, this inevitably sacrificed the crystallinity of the fillers due to the intense shear force introduced by the ball milling process. Thus, it was desirable to develop an effective method to promote the dispersion of fillers in the matrix as well as maintain the quality of the fillers.

Conflicts of interest

There are no conflicts of interest to declare.

Acknowledgements

This work was supported by the National Natural Science Foundation of China (No. 51522211, 51602339 and 51703241), the Key Research Program of Frontier Science of Chinese Academy of Sciences (No. QYZDB-SSW-SLH031), the Thousand Youth Talents Plan, the Postdoctoral Foundation of China (No. 2016M601905 and 2017M621855), the Natural Science Foundation of Jiangsu Province, China (No. BK20160399), the Postdoctoral Foundation of Jiangsu Province (No. 1601065B), and the Science and Technology Project of Suzhou, China (No. SZZS201508).
References

1 D. Lu and C. P. Wong, Materials for advanced packaging, Springer, New York, 2009.

2 C. P. Wong, K. S. Moon and Y. Li, Nano-bio-electronic, photonic and MEMS packaging, Springer, New York, 2010.

3 A. L. Moore and L. Shi, Mater. Today, 2014, 17, 163–174.

4 Y. X. Wang, N. Xu, D. Y. Li and J. Zhu, Adv. Funct. Mater., 2017, 27, 1604134.

5 A. A. Balandin, S. Ghosh, W. Z. Bao, I. Calizo, D. Teweldebrhan, F. Miao and C. N. Lau, Nano Lett., 2008, 8, 902–907.

6 H. S. Kim, J. H. Kim, W. Y. Kim, H. S. Lee, S. Y. Kim and M. S. Khil, Carbon, 2017, 119, 40–46.

7 Z. Luo, J. Maassen, Y. X. Deng, Y. C. Du, R. P. Garrels, M. S. Lundstrom, P. D. Ye and X. F. Xu, Nat. Commun., 2015, 5, 8572.

8 C. R. Dean, A. F. Young, I. Meric, C. Lee, L. Wang, S. Sorgenfrei, K. Watanabe, T. Taniguchi, P. Kim, K. L. Shepard and J. Hone, Nat. Nanotechnol., 2010, 5, 722–726.

9 C. Zhi, Y. Bando, T. Terao, C. Tang, H. Kuwahara and D. Golberg, Adv. Funct. Mater., 2009, 19, 1857–1862.

10 K. C. Yung and H. Liem, J. Appl. Polym. Sci., 2007, 106, 3587–3591.

11 Y. Xu and D. D. L. Chung, Compos. Interfaces, 2000, 7, 243–256.

12 Z. Lin, Y. Liu, S. Raghavan, K. S. Sitaraman and C. P. Wong, ACS Appl. Mater. Interfaces, 2013, 5, 7633–7640.

13 H. S. Lim, J. W. Oh, S. Y. Kim, M. J. Yoo, S. D. Park and W. S. Lee, Chem. Mater., 2013, 25, 3315–3319.

14 Z. Kuang, Y. Chen, Y. Lu, L. Liu, S. Hu, S. Wen, Y. Mao and L. Zhang, Small, 2015, 11, 1655–1659.

15 D. Golberg, Y. Bando, Y. Huang, T. Terao, M. Mitome, C. Tang and C. Zhi, ACS Nano, 2010, 4, 2979–2993.

16 G. R. Bhimanapati, N. R. Glavin and J. A. Robinson, Semicond. Semimetals, 2016, 95, 101–147.

17 W. L. Song, P. Wang, L. Cao, A. Anderson, M. J. Meziani, A. J. Farr and Y. P. Sun, Angew. Chem., Int. Ed., 2012, 51, 6498–6501.

18 L. Boldrin, F. Scarpa, R. Chowdhury and S. Adhikari, Nanotechnology, 2011, 22, 505702.

19 K. Rajesh, G. Rajasekaran and P. Avinash, Nanotechnology, 2016, 27, 085706.

20 J. C. Meyer, A. Chuvilin, G. Algarsa-Siller, J. Biskupek and U. Kaiser, Nano Lett., 2009, 9, 2683–2689.

21 R. V. Gorbachev, I. Riaz, R. R. Nair, R. Jalil, L. Britnell, B. D. Belle, E. W. Hill, K. S. Novoselov, K. Watanabe, T. Taniguchi, A. K. Geim and P. Blake, Small, 2011, 7, 465–468.

22 D. Pacilé, J. C. Meyer, C. Ö. Girit and A. Zettl, Appl. Phys. Lett., 2008, 92, 131307.

23 L. H. Li, Y. Chen, G. Behan, H. Zhang, M. Petrovich and A. M. Glushenkov, J. Mater. Chem., 2011, 21, 11862.

24 Deepika, L. H. Li, A. M. Glushenkov, S. K. Hait, P. Hodgson and Y. Chen, Sci. Rep., 2014, 4, 7288.

25 D. Fan, J. Feng, J. Liu, T. Gao, Z. Ye, M. Chen and X. Lv, Ceram. Int., 2016, 42, 7155–7163.

26 Y. Yao, Z. Lin, Z. Li, X. Song, K. S. Moon and C. P. Wong, J. Mater. Chem., 2012, 22, 13494.

27 Z. Lin, A. McNamara, Y. Liu, K. S. Moon and C. P. Wong, Compos. Sci. Technol., 2014, 90, 123–128.

28 L. Lin, X. Zhitao, H. Daqiang, W. Guotao, L. Bin and C. Ping, Chem. Lett., 2013, 42, 1415–1416.

29 W. Q. Han, L. Wu, Y. Zhu, K. Watanabe and T. Taniguchi, Appl. Phys. Lett., 2008, 93, 223103.

30 Y. Lin, T. V. Williams, T. B. Xu, W. Cao, H. E. Elsayed-Ali and J. W. Connell, J. Phys. Chem. C, 2011, 115, 2679–2685.

31 R. J. Smith, P. J. King, M. Lotya, C. Wirtz, U. Khan, S. De, A. O’Neill, G. S. Duesberg, J. C. Grunlan, G. Moriarty, J. Chen, J. Wang, A. I. Minett, V. Nicolosi and J. N. Coleman, Adv. Mater., 2011, 23, 3944–3948.

32 C. Zhi, Y. Bando, C. Tang, H. Kuwahara and D. Golberg, Adv. Mater., 2009, 21, 2889–2893.

33 Y. Wang, Z. Shi and J. Yin, J. Mater. Chem., 2011, 21, 11371–11377.

34 J. N. Coleman, M. Lotya, A. O’Neill, S. D. Bergin, P. J. King, U. Khan, K. Young, A. Gaucher, S. De, R. J. Smith, I. V. Shvets, S. K. Arora, G. Stanton, H. Y. Kim, K. Lee, G. T. Kim, G. S. Duesberg, T. Hallam, J. J. Boland, J. J. Wang, J. F. Donegan, J. C. Grunlan, G. Moriarty, A. Shmeliov, R. J. Nicholls, J. M. Perkins, E. M. Grieveson, K. Theuwissen, D. W. McComb, P. D. Nellist and V. Nicolosi, Science, 2011, 331, 568–571.

35 K. G. Zhou, N. N. Mao, H. X. Wang, Y. Peng and H. L. Zhang, Angew. Chem., Int. Ed., 2011, 50, 10839–10842.

36 L. Cao, S. Emami and K. L. Dadf, Mater. Express, 2014, 4, 165–171.

37 H. B. Cho, T. Nakayama, H. Suematsu, T. Suzuki, W. Jiang, K. Niihara, E. Song, N. S. A. Eom, S. Kim and Y. H. Choa, Compos. Sci. Technol., 2016, 129, 205–213.

38 M. T. Paffett, R. J. Simonson, P. Papin and R. T. Paine, Surf. Sci., 1990, 232, 286–296.

39 A. Nagashima, N. Tejima, Y. Gamou, T. Kawai and C. Oshima, Phys. Rev. B, 1995, 51, 4606–4613.

40 F. Müller, K. Stöwe and H. Sachdev, Chem. Mater., 2005, 17, 3464–3467.

41 W. Auwärter, H. U. Suter, H. Sachdev and T. Greber, Chem. Mater., 2004, 16, 343–345.

42 M. Corso, W. Auwärter, M. Muntwiler, A. Tamai, T. Greber and J. Osterwalder, Science, 2004, 303, 217–220.

43 N. A. Vinogradov, A. A. Zakharov, M. L. Ng, A. Mikkelsen, E. Lundgren, N. Märtenson and A. B. Preobrajenski, Langmuir, 2012, 28, 1775–1781.

44 F. Müller, S. Hufner and H. Sachdev, Surf. Sci., 2008, 602, 3467–3476.

45 M. P. Allan, S. Berner, M. Corso, T. Greber and J. Osterwalder, Nanoscale Res. Lett., 2007, 2, 94.

46 A. B. Preobrajenski, M. A. Nesterov, M. L. Ng, A. S. Vinogradov and N. Märtenson, Chem. Phys. Lett., 2007, 446, 119–123.

47 A. B. Preobrajenski, A. S. Vinogradov and N. Märtenson, Surf. Sci., 2005, 582, 21–30.
93 H. J. Ahn, Y. J. Eoh, S. D. Park and E. S. Kim, *Thermochim. Acta*, 2014, [590](590), 138–144.

94 S. Takahashi, Y. Imai, A. Kan, Y. Hotta and H. Ogawa, *J. Alloys Compd.*, 2014, [615](615), 141–145.

95 R. M. Erb, R. Libanori, N. Rothfuchs and A. R. Studart, *Science*, 2012, [335](335), 199–204.

96 H. B. Cho, Y. Tokoi, S. Tanaka, H. Suematsu, T. Suzuki, W. Jiang, K. Niihara and T. Nakayama, *Compos. Sci. Technol.*, 2011, [71](71), 1046–1052.

97 K. Kim and J. Kim, *Int. J. Therm. Sci.*, 2016, [100](100), 29–36.

98 C. Yuan, B. Duan, L. Li, B. Xie, M. Huang and X. Luo, *ACS Appl. Mater. Interfaces*, 2015, [7](7), 13000–13006.

99 H. B. Cho, T. Nakayama, Y. Tokoi, S. Endo, S. Tanaka, T. Suzuki, W. Jiang, H. Suematsu and K. Niihara, *Compos. Sci. Technol.*, 2010, [70](70), 1681–1686.

100 T. Fujihara, H. B. Cho, T. Nakayama, T. Suzuki, W. Jiang, H. Suematsu, H. D. Kim, K. Niihara and J. Blendell, *J. Am. Ceram. Soc.*, 2012, [95](95), 369–373.

101 H. Zhu, Y. Li, Z. Fang, J. Xu, F. Cao, J. Wan, C. Preston, B. Yang and L. Hu, *ACS Nano*, 2014, [8](8), 3606–3613.

102 K. Wu, J. C. Fang, J. R. Ma, R. Huang, S. G. Chai, F. Chen and Q. Fu, *ACS Appl. Mater. Interfaces*, 2017, [9](9), 30035–30045.

103 J. C. Zheng, L. Zhang, A. V. Kretinin, S. V. Morozov, Y. B. Wang, T. Wang, X. Li, F. Ren, J. Zhang and C. Y. Lu, *2D Mater.*, 2016, [3](3), 011004.

104 L. Fu, T. Wang, J. Yu, W. Dai, H. Sun, Z. Liu, R. Sun, N. Jiang, A. Yu and C. T. Lin, *2D Mater.*, 2017, [4](4), 025047.

105 X. Zeng, Y. Yao, Z. Gong, F. Wang, R. Sun, J. Xu and C. P. Wong, *Small*, 2015, [11](11), 6205–6213.

106 J. Chen, X. Huang, Y. Zhu and P. Jiang, *Adv. Funct. Mater.*, 2017, [27](27), 1604754.

107 J. Yang, L. S. Tang, R. Y. Bao, L. Bai, Z. Y. Liu, W. Yang, B. H. Xie and M. B. Yang, *J. Mater. Chem. A*, 2016, [4](4), 18841–18851.

108 B. H. Xie, X. Huang and G. J. Zhang, *Compos. Sci. Technol.*, 2013, [85](85), 98–103.

109 Y. Xue, X. Jin, Y. Fan, R. Tian, X. Xu, J. Li, J. Lin, J. Zhang, L. Hu and C. Tang, *Polym. Compos.*, 2014, [35](35), 1707–1715.

110 K. Kim and J. Kim, *Ceram. Int.*, 2014, [40](40), 5181–5189.

111 Z. Wang, Y. Fu, W. Meng and C. Zhi, *Nanoscale Res. Lett.*, 2014, [9](9), 643.

112 X. J. Wang, L. Z. Zhang and L. X. Pei, *J. Appl. Polym. Sci.*, 2014, [131](131), 39550.

113 T. L. Li and S. L. Hsu, *J. Phys. Chem. B*, 2010, [114](114), 6825–6829.

114 K. C. Yung and H. Liem, *J. Appl. Polym. Sci.*, 2007, [106](106), 3587–3591.

115 W. Y. Zhou, S. H. Qi, H. Z. Zhao and N. L. Liu, *Polym. Compos.*, 2007, [28](28), 23–28.

116 T. L. Li and S. L. C. Hsu, *J. Phys. Chem. B*, 2010, [114](114), 6825–6829.

117 S. Kemaloglu, G. Ozkoc and A. Aytaç, *Thermochim. Acta*, 2010, [499](499), 40–47.

118 L. Fang, C. Wu, R. Qian, L. Xie, K. Yang and P. Jiang, *RSC Adv.*, 2014, [4](4), 21010–21017.

119 A. Permal, M. Devarajan, H. L. Hung, T. Zahner, D. Lacey and K. Ibrahim, *J. Mater. Sci.*, 2016, [51](51), 7415–7426.

120 Y. X. Fu, Z. X. He, D. C. Mo and S. S. Lu, *Appl. Therm. Eng.*, 2014, [66](66), 493–498.

121 F. Wang, X. Zeng, Y. Yao, R. Sun, J. Xu and C. P. Wong, *Sci. Rep.*, 2016, [6](6), 19394.

122 Y. Yao, X. Zeng, R. Sun, J. B. Xu and C. P. Wong, *ACS Appl. Mater. Interfaces*, 2016, [8](8), 15645–15653.

123 M. H. Tsai, I. H. Tseng, J. C. Chiang and J. J. Li, *J. Appl. Polym. Sci.*, 2016, [120](120), 8639–8645.

124 X. Cui, P. Ding, N. Zhuang, L. Shi, N. Song and S. Tang, *ACS Appl. Mater. Interfaces*, 2015, [7](7), 19068–19075.

125 Y. Xiao, W. Wang, T. Lin, X. Chen, Y. Zhang, J. Yang, Y. Wang and Z. Zhou, *J. Phys. Chem. C*, 2016, [120](120), 6344–6355.