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Controlling the electrical conductive network formation in nanorod filled polymer nanocomposites by tuning nanorod stiffness†

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In this work, by employing a coarse-grained molecular dynamics simulation, we have investigated the effect of the nanorod (NR) stiffness on the relationship between the NR microstructure and the conductive probability of NR filled polymer nanocomposites (PNCs) under the quiescent state and under the shear field. The conductive probability of PNCs is gradually enhanced with the increase of NR stiffness in the quiescent state; however, it first increases and then decreases under the shear field. As a result, the largest conductive probability appears at moderate NR stiffness, which results from the competition between the improved effective aspect ratio of the NR and the breakage of the conductive network. Meanwhile, compared with in the quiescent state, under the shear field the decrease or the increase of the conductive probability depends on the NR stiffness. At low NR stiffness, the increase of the effective aspect ratio of NR enhances the conductive probability, while at high NR stiffness, the breakage of the conductive network reduces the conductive probability. For flexible NRs, the conductive probability first increases and then decreases with increasing the shear rate. The maximum effective aspect ratio of NRs appears at the moderate shear rate, which is consistent with the conductive probability. In summary, this work presents some further understanding about how NR stiffness affects the electric conductive properties of PNCs under the shear field.

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1. Introduction

Addition of a conductive filler (such as carbon black (CB), carbon nanotubes (CNTs), graphene) into an insulating polymer matrix can markedly improve the electrical properties of polymer nanocomposites (PNCs) by orders of magnitude.^{1–5} The conductive network is formed if the filler volume fraction reaches the percolation threshold, which can be described by the percolation theory.⁶ From the application viewpoint, it is very important to reduce the percolation threshold, which can improve the electrical conductivity and reduce cost for manufacturing materials.

Currently, some reviews^{2,7,8} have summarized the methods to tune and control the conductive network, which can help

understand the relationship between the experimental parameters and the conductivity. For example, the polymer–filler interaction affects the formation of the conductive network, which further influences the electrical properties.^{9,10} The dependence of the percolation threshold on the aspect ratio of CNT can be described by a model, which could optimize the CNTs' processing conditions.¹¹ The shear is an important method during the processing of PNCs, which will affect the electrical conductivity. To correctly predict it, it is very important to understand the response of the conductive network under the shear field. Kharchenko *et al.*¹² first reported the destruction of the conductive network under the shear field for the initially partially aggregated CNTs in PP matrix. The shear field can destroy the conductive network, which reduce the electrical properties.¹³ For the well dispersed CNTs, the insulator–conductor transition happens, which is attributed to the formation of the conductive network induced by the shear field.¹⁴ However, it is very difficult to intuitively analyze the variation of the conductive network and electrical properties by experimental methods. Computer simulation and models provide another good choice to study the electrical conductivity behavior. Currently, by adopting the critical path method, the electrical conductivity can be extracted from the critical distance between nanorods (NRs).^{15–18} By employing the continuum percolation theory, the percolation threshold is

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reported to depend on the aspect ratio¹⁹ and the size polydispersity^{20,21} of NR. The depletion effect of insulating spheres (not conductive filler) can induce the effective attraction between NRs (conductive filler), which enhance the conductivity of PNCs.^{22,23} By utilizing Monte Carlo model, it reveals an exponential relationship between the fiber aspect ratio and the percolation threshold.²⁴ In addition, the high size polydisperse increases the percolation threshold, which is approximately inversely upon the weight-averaged aspect ratio.²⁵ The contribution of longer CNTs to the formation of conductive network is through linking the shorter CNTs.^{26,27} By employing molecular dynamics simulation, nanorods are more effective in forming a percolating network at a lower volume fraction than nanoplates and nanospheres.²⁸ The anisotropy of the electrical property of aligned CNT/polymer composites is mainly affected by the average conductive pathway density of CNTs.²⁹ Under the shear field, the shear-induced CNT orientation reduces the percolation threshold of PNCs along the shear direction.³⁰ Especially, the highest electrical conductivity occurs when the CNTs are partially rather than perfectly aligned.^{31,32} Interestingly, when the shear rate is beyond a critical value, the conductive network is broken down because of CNT aggregation.³³ In addition, both the shear rate and the CNT aspect ratio influence the formation and destruction of the conductive aggregates, which further affect the electrical conductivity.^{34,35}

In fact, fillers are flexible if they have a very large aspect ratio.^{36,37} The effective aspect ratio of flexible fillers is lower than that of the straight fillers, which will have a considerable effect on the electrical conductivity. It is noted that the effective aspect ratio is defined as the ratio of the end-to-end distance of filler to the diameter of filler (1σ). Current works^{26,28,38–40} indicate that the flexible fillers tend to increase the percolation threshold. However, the effect of filler stiffness on the conductive behavior of PNCs under the shear field has not been investigated to our knowledge. In this work, by adopting a coarse-grain model, first, the microstructure and the kinetics of the NR in the matrix is characterized for different NR stiffness. Then, the conductive probability of PNCs is calculated under the quiescent state and under the shear field. Here, we mainly considered three kinds of conductive probabilities, namely homogeneous conductive probability A , directional conductive probability A_{\parallel} parallel to the shear field direction, and directional conductive probability A_{\perp} perpendicular to the shear field direction.

2. Simulation details

In this work, a coarse-grained model of NRs filled polymer nanocomposites is adopted. The classic bead-spring model⁴¹ is adopted to simulate polymer chains which consist of thirty beads. The total number of polymer beads for each system is fixed at 42 000. Each NR contains ten beads. Two symbols σ and m are adopted to stand for the diameter and mass of each bead (polymer and NRs), respectively.^{42,43} Although these chains are rather short compared to real chains, they are already able to display the static and dynamic characteristic behavior of long chains. Each bond in this model corresponds to three to six

covalent bonds along the backbone of a real chemical chain when mapping the coarse-grained model to a real polymer.

A stiff finite extensible nonlinear elastic (FENE) potential⁴⁴ is used to represent the bonded interaction between the adjacent beads including both polymer chains and NRs, which is given by

$$V_{\text{FENE}} = -0.5kR_0^2 \ln \left[1 - \left(\frac{r}{R_0} \right)^2 \right] \quad (1)$$

where $k = 30 \frac{\varepsilon}{\sigma^2}$ and $R_0 = 1.5\sigma$, guaranteeing a certain stiffness of the bonds while avoiding high-frequency modes and chain crossing.

The non-bonded interactions between all the beads are described using a truncated and shifted Lennard-Jones (LJ) potential, given by

$$U(r) = \begin{cases} 4\varepsilon \left[\left(\frac{\sigma}{r} \right)^{12} - \left(\frac{\sigma}{r} \right)^6 \right] & r < r_{\text{cutoff}} \\ 0 & r \geq r_{\text{cutoff}} \end{cases} \quad (2)$$

where r_{cutoff} stands for the distance at which the interaction is truncated and shifted so that the potential is continuous at $r = r_{\text{cutoff}}$. The polymer–polymer interaction parameter and its cutoff distance are $\varepsilon_{\text{pp}} = 1.0$ and $r_{\text{pp}} = 2 \times 2^{1/6}$, and the NR–NR interaction parameter and its cutoff distance are set to be $\varepsilon_{\text{nn}} = 1.0$ and $r_{\text{nn}} = 1.12$. The polymer–NR interaction parameter and its cutoff distance are $\varepsilon_{\text{np}} = 1.0$ and $r_{\text{np}} = 2 \times 2^{1/6}$, which simulates the weak attractive interaction.

The rod-like character of the NR is enforced by a bending potential, given by

$$U_{\text{angle}} = K_{\text{stiffness}}(\theta - \theta_0)^2 \quad (3)$$

where θ is the bending angle formed by three consecutive rod beads, θ_0 is set to be 180, and $K_{\text{stiffness}}$ is changed from 0 to 100 to simulate different NR stiffness.

Since it is not our aim to study a specific polymer, the LJ units (distance = σ , mass = m , interaction energy = ε) are set to unit. The reduced units of other quantities (temperature, density, pressure and time) are adopted, defined as: $k_{\text{B}}T/\varepsilon$, $\rho\sigma^3$, $P\sigma^3/\varepsilon$ and t/τ ($\tau = \sigma\sqrt{m/\varepsilon}$), respectively. Similar to our previous work,^{44–46} first, all the polymer chains and NRs are put into a large box. Then we adopted the NPT ensemble to compress the system for 20000 τ , where the temperature and pressure are fixed at $T^* = 1.0$ and $P^* = 0.0$ respectively by using the Nose–Hoover temperature thermostat and pressure barostat. Further equilibration under NVT with $T^* = 1.0$ is performed for 50000 τ . It is noted that each chain has moved at least $2R_{\text{g}}$ during the long simulation process. We have checked that polymer chains and NRs have experienced fully relaxed, reaching their equilibrated states for all systems. The equilibrium number density of polymer beads reaches nearly 0.85 corresponding to the density of polymer melts. Such equilibrated structures are then used as starting structures for production runs of the structural and dynamical analysis. During the simulation process, periodic boundary conditions are employed in all three directions. The velocity-Verlet algorithm is applied



to integrate the equations of motion with a timestep of $\delta t = 0.001\tau$, where the time is reduced by LJ unit time (τ). All MD runs are carried out by using the large scale atomic/molecular massively parallel simulator (LAMMPS).⁴⁷

First, to determine whether the conductive network is formed, a criterion is used to check whether any two NRs are connected. If the gap of two NRs is less than the tunneling distance (TD), they are considered to be connected. TD can stand for the contact conductance between a pair of the NRs.⁴⁸ High TD indicates the high contact conductance. Here, the tunneling distance is chosen to be 1.0σ from the two aspects (one is the sandwiched polymer chain, another is that the ratio of CNT diameter to the maximum TD is nearly 1.0 in experiment⁴⁹). It is noted that TD impacts upon the percolation threshold, which scales with the reciprocal TD for long rods. However, it does not affect the trend of the results. In addition, TD is fixed even though it maybe affected by both shear and bending flexibility.

At the beginning of the computational implementation, each NR is assigned a site number and a cluster number.^{24,50} The site number is equal to the cluster number, ranging from 1 to N , where N is the total number of NRs. Then each NR is checked for connection with others. If two NRs are connected, they will be assigned a common cluster number. Finally, all the NRs with the same cluster number are in the same cluster. As a result, NRs with different cluster numbers are not connected. If the NR network spans one direction continuously from one side to another, the system is conductive in this direction. If the NR network spans the three-dimensional directions continuously, the system is homogeneously conductive in three directions. Last, more than 10 000 equilibrated configurations are dumped to perform the data analysis. The time interval between two continuous frames is 10τ . Finally, the number of the configurations, which is conductive in the three-dimensional directions or in one direction, is counted. The conductive probability \mathcal{A} , directional conductive probability \mathcal{A}_{\parallel} parallel to the shear field direction, and directional conductive probability \mathcal{A}_{\perp} perpendicular to the shear direction are used to represent the formation probability of the filler network, which spans the systems in the three-dimensional directions, parallel to the shear direction, and perpendicular to the shear direction, respectively.

3. Results and discussion

3.1 Conductive property of PNCs

3.1.1 Effect of the nanorod stiffness $K_{\text{stiffness}}$. Stiffness is a prevailing feature of fillers in PNCs. Flexible fillers (low stiffness) tend to have a lower effective aspect ratio than the flexible fillers (high stiffness). Therefore, filler stiffness will have a considerable effect on the electrical conductivity by using a percolation model.³⁹ The aggregation structure and the rotational diffusion of NR with different stiffness in the polymer matrix has been discussed in the ESI.† The conductive probability \mathcal{A} as a function of the NR volume fraction (ϕ) for different NR stiffness is presented in Fig. 1. It is found that variation of \mathcal{A} with ϕ shows a typical percolation phenomenon. In the percolated region, the conductive network is gradually formed and

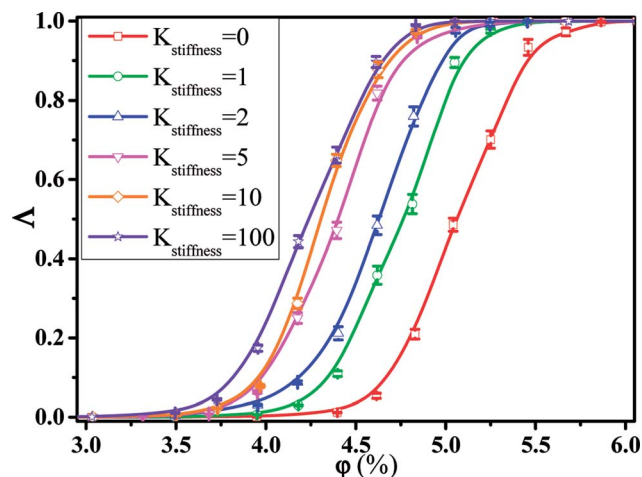


Fig. 1 Conductive probability \mathcal{A} of polymer nanocomposites as a function of the nanorod volume fraction ϕ for different nanorod stiffness ($K_{\text{stiffness}}$). ($T^* = 1.0$).

developed. As a result, the NR conductive network is strongly dependent on the NR volume fraction, which leads to the quick increase of \mathcal{A} with tiny variation of ϕ . Here, we defined the percolation threshold ϕ_c as the NR volume fraction at $\mathcal{A} = 0.5$,⁵¹ which gradually decreases from 5.06%, 4.77%, 4.63%, 4.41%, 4.31% to 4.23% with the increase of $K_{\text{stiffness}}$ from 0, 1, 2, 5, 10 to 100, respectively. The reason for the continuous decrease of ϕ_c is that NRs with high stiffness gradually straighten, which leads to the large effective aspect ratio. High effective aspect ratio indicates the straight NR, which benefits the formation of the continuous NR conductive network throughout the systems. According to the excluded volume theory,^{36,52} NRs with high effective aspect ratio can be easily connected with other NRs to form a conductive network, leading to a low percolation threshold. This is consistent with our results. To characterize the NR conductive network, we introduced two parameters, the main cluster size C_n (the number of NR beads within the main cluster), and the total number of clusters N_c to analyze the conductive probability.⁴⁵ Generally, at the same NR volume fraction, the system with higher C_n and smaller N_c has more opportunity to form the conductive network. As shown in Fig. 2, in the percolation region, it is found that C_n decreases and N_c increases with increasing NR stiffness from 0 to 100, which is consistent with \mathcal{A} . The difference of N_c is small for $K_{\text{stiffness}} \geq 5$, which maybe attributed to the similar C_n and \mathcal{A} . In addition, some typical snapshots of the main cluster at different NR stiffness and NR volume fractions are shown in Fig. 3. It presents that at low ϕ , the main cluster can not span the system continuously. With the increase of ϕ , the main cluster size gradually increases, where a continuous NR network forms through the systems. These results further confirm that the high NR stiffness leads to high effective aspect ratio, which can help to form the conductive network.

3.1.2 Effect of the external field. The external shear field is a very common means to manufacture materials.^{7,53} When the materials are exposed to the shear field, the conductive network is broken down, which affects the conductive



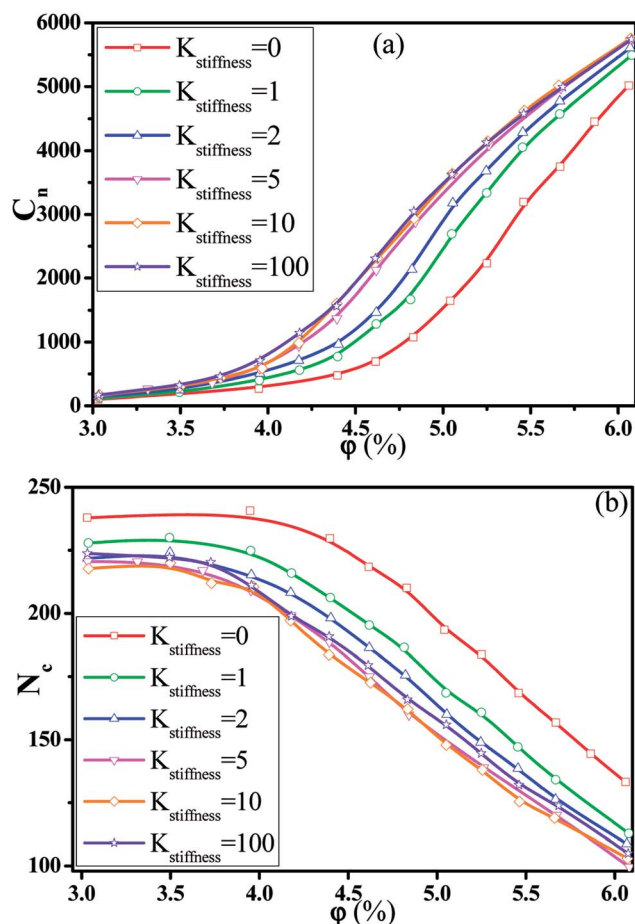


Fig. 2 Change of (a) the main cluster size C_n and (b) the total number of clusters N_c as a function of the nanorod volume fraction ϕ for different nanorod stiffness ($K_{\text{stiffness}}$). ($T^* = 1.0$).

probability. After the shear field is stable, a new network is formed. Here, the SLLOD equations⁵⁴ is adopted to realize the shear simulation, which is one of the most widely used methods to study the shearing systems. We used the special Lees-Edwards “sliding brick” boundary conditions⁵⁵ for the SLLOD method, which is very effective for the shear simulation. The shear field is exerted on the simulation box by moving top xy plane of the simulation box along the x direction. Here, the orientation of NR is used to determine whether the shear flow is stable.

First, we investigated the effect of the shear rate $\dot{\gamma}$ on the conductive probability of PNCs by fixing the NR $K_{\text{stiffness}}$ equal to 2, where NRs can straighten under the shear field. Change of A , A_{\parallel} , and A_{\perp} with different NR volume fractions and shear rates under a stable shear flow is presented in Fig. 4. It is noted that two directional conductive probabilities along the gradient direction and the vorticity direction are averaged as the directional conductive probability perpendicular to the shear direction A_{\perp} because they are very similar. It is interesting to find that A first increases and then decreases with increasing $\dot{\gamma}$, which can be clearly reflected by the change of ϕ_c in Fig. 5(a). It is noted that A depends on both A_{\parallel} and A_{\perp} . The continuous increase of A_{\parallel} is generally attributed to the NR orientation along

the shear direction. A_{\perp} shows a similar trend with A as the increase of $\dot{\gamma}$ even though the difference of A_{\perp} is small for different $K_{\text{stiffness}}$. To further explore it, first the second-order Legendre polynomials $\langle P_2 \rangle$ is used to denote the NR orientation degree at different shear rates, given by

$$\langle P_2 \rangle = (3\langle \cos^2 \theta_2 \rangle - 1)/2 \quad (4)$$

where θ_2 denotes the angle between the end-to-end vector of NRs and the shear direction. From Fig. 5(b), the results present that NRs gradually orientate along the shear direction. Then, we calculated the end-to-end distance of NR $|\vec{R}_{ee}| = |\vec{r}_1 - \vec{r}_n|$, where \vec{r}_1 and \vec{r}_n are the position vector of the first and the last beads of NR in Fig. 5(a). It is interesting to find that $|\vec{R}_{ee}|$ (which can reflect the effective aspect ratio of NR) first gradually increases and then decreases with $\dot{\gamma}$. The extremum of ϕ_c and $|\vec{R}_{ee}|$ appears at the same $\dot{\gamma}$. Last, we characterized the NR dispersion state under the shear field by calculating the inter-nanorod radial distribution function (RDF), which is shown in Fig. S4(a).† The peaks at $r = 1\sigma$, 2σ and 3σ gradually increase with $\dot{\gamma}$, which reflects that more NRs tend to aggregate directly.

According to the results mentioned above, in the quiescent state NRs disperse randomly in the matrix, leading to a uniform conductivity probability along three different directions. Under the shear field, on one hand, NRs begin to orientate along the shear field; on the other hand, more NRs tend to aggregate directly. They are responsible for the increase of A_{\parallel} with $\dot{\gamma}$ in Fig. 4(b). NRs with low stiffness ($K_{\text{stiffness}} = 2$) first straighten with $\dot{\gamma}$; however, it does not have enough time to further straighten at $\dot{\gamma} \geq 0.1$, which leads to the decrease of R_{ee} in Fig. 5(a). As a result, the maximum effective aspect ratio of NRs appears at $\dot{\gamma} = 0.05$. In general, the NR orientation destroys the inter-nanorod connection perpendicular to the shear direction, which reduces A_{\perp} . However, at $\dot{\gamma} < 0.1$, the increase of A_{\perp} by the improved effective aspect ratio of NRs makes up the decrease of A_{\perp} by the NR orientation, which eventually leads to the weak increase of A_{\perp} . At $\dot{\gamma} \geq 0.1$, A_{\perp} decreases again because of the reduced effective aspect ratio of NRs in Fig. 5(a). In summary, first A strongly depends on A_{\perp} , rather than A_{\parallel} . This is because the conductive network parallel to the shear direction is much easier to be formed than that perpendicular to the shear direction. To better understand it, we also calculated the main cluster size C_n to analyze the conductive network in Fig. S4(b).† The results show that C_n first increases and then decreases with $\dot{\gamma}$, which is consistent with A . To observe the NR conductive network, we presented the snapshots of some typical systems ($\phi = 4.40\%$) with different $\dot{\gamma}$ in Fig. S5.† It presents that NRs gradually align along the shear direction. Meanwhile, the main cluster is gradually broken down, which is reflected by the decrease in the number of red beads in Fig. S5.† In total, our results show that the original NR conductive network is broken down and a new one is formed. Accompany with it, the effective aspect ratio and the orientation of NRs along the shear direction changes with $\dot{\gamma}$, which synergistically affects the conductive probability.



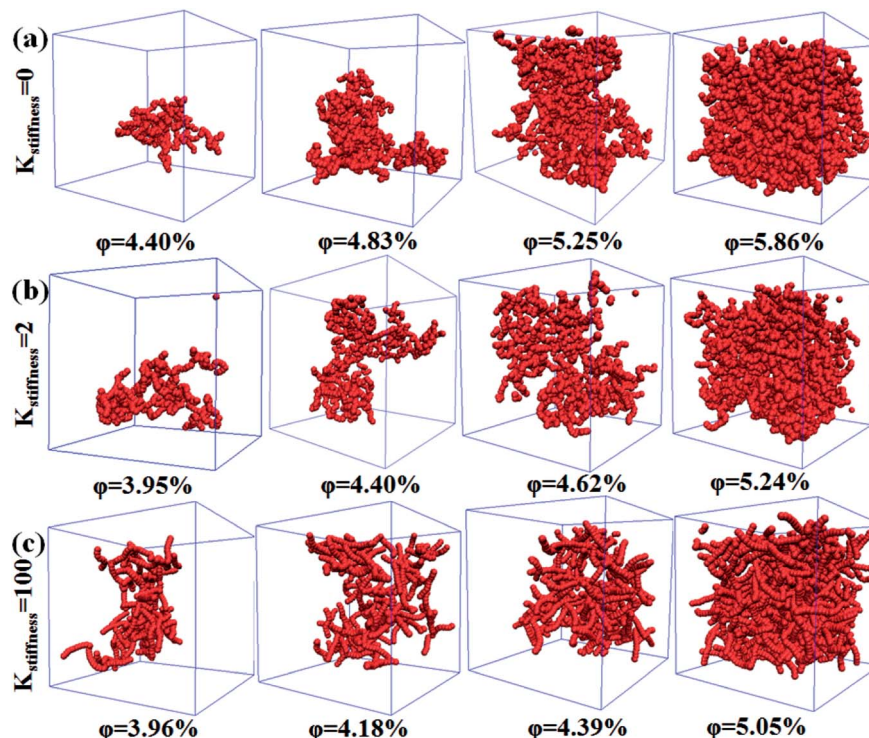


Fig. 3 Some typical snapshots of the main cluster with different nanorod stiffness (a) $K_{\text{stiffness}} = 0$, (b) $K_{\text{stiffness}} = 2$, and (c) $K_{\text{stiffness}} = 100$ at four nanorod volume fractions ϕ . The red spheres denote the nanorods with the main cluster. ($T^* = 1.0$).

Then, we investigated the effect of the NR stiffness ($K_{\text{stiffness}}$) on the conductive probability of PNCs by fixing the shear rate $\dot{\gamma} = 0.1$. Fig. 6 shows variation of Λ , Λ_{\parallel} , and Λ_{\perp} for different NR volume fractions and $K_{\text{stiffness}}$ in a steady shear flow. Under the shear field, Λ in Fig. 6(a) first increases and then decreases with the increase of $K_{\text{stiffness}}$ at $\dot{\gamma} = 0.1$, which is different with that at $\dot{\gamma} = 0.0$ in Fig. 1. To clearly present it, the percolation threshold ϕ_c is shown in Fig. 7(a) for different $K_{\text{stiffness}}$ at $\dot{\gamma} = 0.0$ and 0.1. Similarly, the increase of Λ_{\parallel} in Fig. 6(b) is attributed to the gradual NR orientation with $K_{\text{stiffness}}$ in Fig. S6.† Meanwhile, Λ and Λ_{\perp} have the same trend with $K_{\text{stiffness}}$. Under the shear field, on one hand the breakage of conductive network reduces Λ . On the other hand, the increase of the effective aspect ratio of NR enhances Λ . As shown in Fig. 7(b), at low $K_{\text{stiffness}} < 5$, R_{ee} of NR at $\dot{\gamma} = 0.1$ increases by more than 7% compared with that at $\dot{\gamma} = 0.0$. The increase of Λ by the improved effective aspect ratio of NR can make up the decrease of Λ induced by the breakage of conductive network, which enhances Λ . However, at high $K_{\text{stiffness}} > 5$, both the breakage of the conductive network and the NR stiffness determines the variation in Λ because of the limited increase of R_{ee} ($\sim 1.5\%$ for $K_{\text{stiffness}} = 100$). Thus, the $K_{\text{stiffness}}$ of NR determines the increase or the decrease of the conductive probability under the shear field compared with that the quiescent state. Similarly, the main cluster size C_n is calculated to analyze the conductive network in Fig. S7.† The results show that C_n first increases and then decreases with $K_{\text{stiffness}}$, which is consistent with Λ . To observe the conductive network, the snapshots of some typical systems ($\phi = 4.40\%$) are presented with different $K_{\text{stiffness}}$ at $\dot{\gamma} = 0.0$ and 0.1 in

Fig. S8(a).† It clearly presents that NRs gradually align along the shear direction from $\dot{\gamma} = 0.0$ to 0.1. Meanwhile, at $\dot{\gamma} = 0.1$, C_n increases for $K_{\text{stiffness}} = 0.0$, while it decreases for $K_{\text{stiffness}} = 100$ compared with those at $\dot{\gamma} = 0.0$. This can be reflected by the decrease in the number in red beads. In total, compared with in the quiescent state, the decrease or the increase of the conductive probability strongly depends on the NR stiffness under the shear field.

Under the shear field, the conductive anisotropy $\Lambda_{\parallel}/\Lambda_{\perp}$ is induced by the NR orientation, defined in eqn (4). Here, we chose systems which contain 4000 NRs ($\phi = 3.95\%$), which is near to the percolation threshold. Otherwise, Λ_{\parallel} or Λ_{\perp} approaches 0 or 1 which induces large error. The result in Fig. S8(b)† indicates that the dependence of $\Lambda_{\parallel}/\Lambda_{\perp}$ on the NR orientation $\langle P_2 \rangle$ can be described by a simple empirical formula for different $K_{\text{stiffness}}$, given by:

$$\ln(\Lambda_{\parallel}/\Lambda_{\perp}) = A \times \langle P_2 \rangle \quad (5)$$

where A is a fitted parameter. In summary, under the shear field the NR network is gradually broken down and orientates along the shear direction, inducing the anisotropy of the conductive probability.

3.2 Discussion

When mapping the coarse-grained model to real polymers,⁴¹ the interaction parameter ϵ is about 2.5–3.4 kJ mol^{−1} for the common polymers. In addition, for this coarse-grained model of polymer chains, the persistence length is about 0.676 σ . For



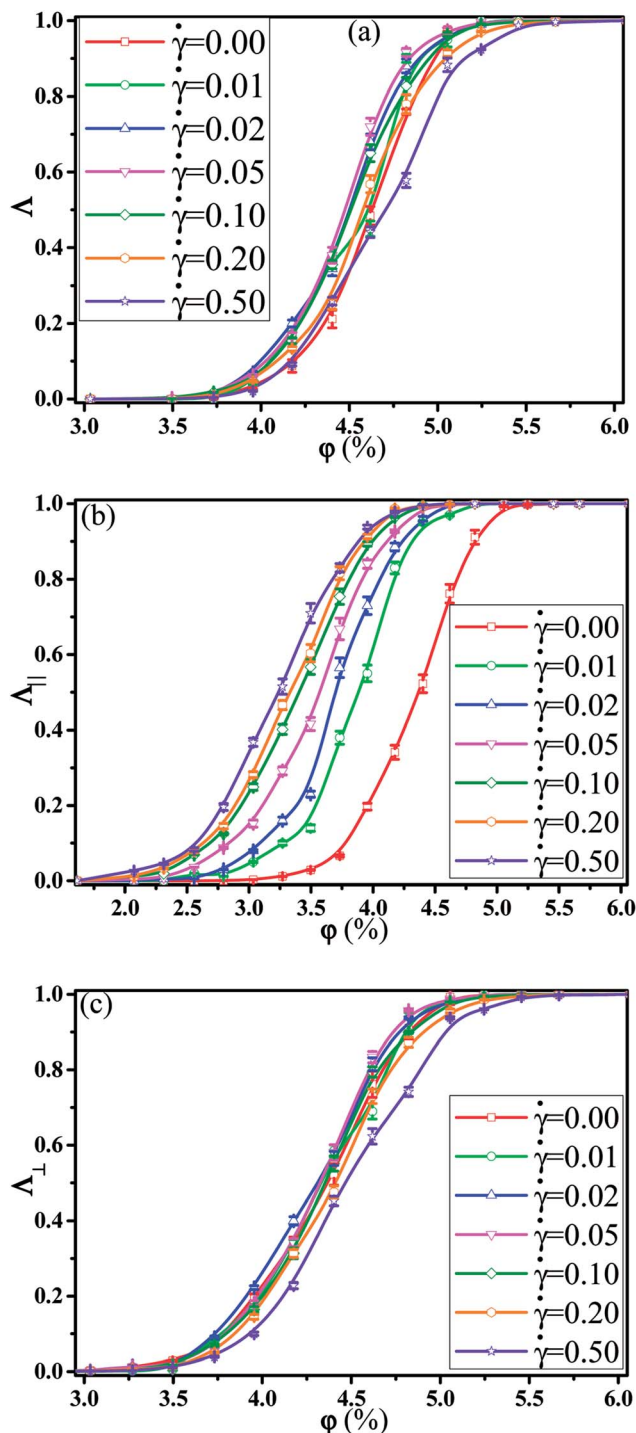


Fig. 4 (a) Homogeneous conductive probability Δ , (b) directional conductive probability $\Delta_{||}$ parallel to the shear direction, and (c) directional conductive probability Δ_{\perp} perpendicular to the shear direction as a function of the nanorod volume fraction ϕ for different shear rates $\dot{\gamma}$. ($T^* = 1.0$, $K_{\text{stiffness}} = 2.0$).

real polymers, the range of the persistence length is between 0.35 nm and 0.76 nm.⁵⁶ Thus, σ can be roughly about 1 nm. Here, polyethylene is used as an example. Each bead with its diameter σ equal to 1 nm roughly corresponds to 5 repeating units which leads to the mass of each bead equal to 140g mol⁻¹. The time scale $\tau = \sigma\sqrt{m/\varepsilon}$ is calculated, which is within the

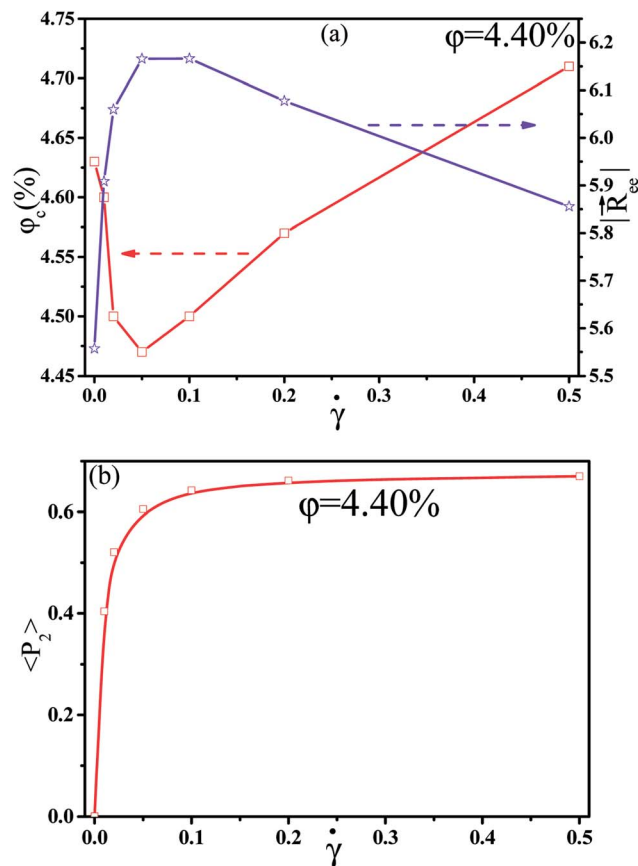


Fig. 5 (a) The left axis denotes the percolation threshold ϕ_c , while the right axis represents the end-to-end distance $|R_{ee}|$ of the nanorod with respect to the shear rate $\dot{\gamma}$. (b) The nanorod orientation $\langle P_2 \rangle$ with respect to the shear rate $\dot{\gamma}$. ($T^* = 1.0$, $\phi = 4.40\%$, $K_{\text{stiffness}} = 2$).

range of about 10 ps. In the ESI,† the rotational diffusion coefficient (D_{rot}) gradually decreases from $15.5 \times 10^{-4} \tau^{-1}$, $12.2 \times 10^{-4} \tau^{-1}$, $9.7 \times 10^{-4} \tau^{-1}$, $7.35 \times 10^{-4} \tau^{-1}$, $5.35 \times 10^{-4} \tau^{-1}$ to $2.43 \times 10^{-4} \tau^{-1}$ with the increase of $K_{\text{stiffness}}$ from 0, 1, 2, 5, 10 to 100, respectively. The shear rate $\dot{\gamma}$ varies from $0.1 \tau^{-1}$ to $0.5 \tau^{-1}$ in

our simulation. Thus, the Peclet number ($Pe = \frac{\dot{\gamma}}{D_{\text{rot}}}$) varies from 64 to 2058, which is comparable with the experimental Peclet numbers.⁵⁷ Thus, our simulation is roughly reasonable. In addition, in experiments because of the attractive interaction between fillers, the fillers always aggregate directly and the conductive network can not be formed. To improve the dispersion state of fillers, an effective strategy is to functionalize the filler surface by grafted polymer chains.^{58–61} As a results, the filler surface is covered with a layer of polymer, which leads to the similar interactions for ε_{pp} , ε_{np} and ε_{nn} . Thus, in this work, ε_{pp} , ε_{np} , and ε_{nn} are set to be 1.0, which roughly can stand for the system with a good dispersion state of grafted fillers in the matrix.

The electrical conductivity is closely dependent on the conductive network, which is mainly affected by several factors, such as filler dispersion,^{23,35} filler shape,^{28,35} filler aspect ratio,^{11,62} filler–filler and polymer–filler interactions,^{10,30} filler alignment,^{29,32} etc. Due to their very large aspect ratio, carbon



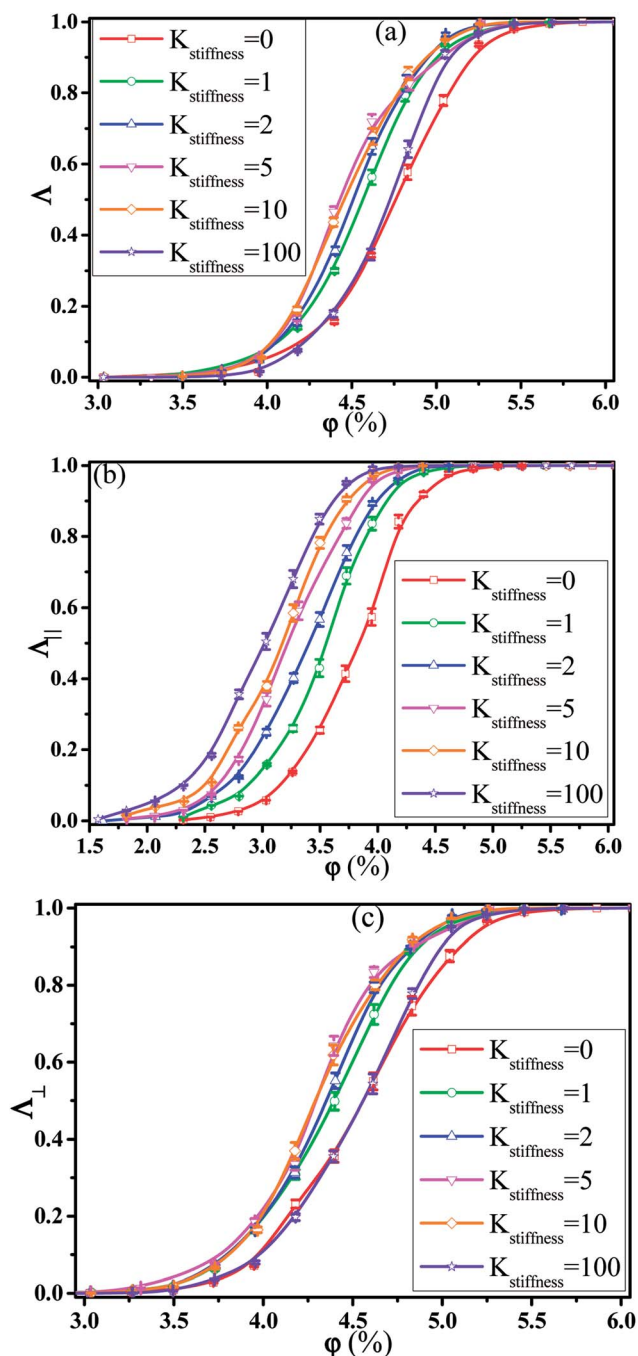


Fig. 6 (a) Homogeneous conductive probability Δ , (b) directional conductive probability Δ_{\parallel} parallel to the shear direction, and (c) directional conductive probability Δ_{\perp} perpendicular to the shear direction as a function of the nanorod volume fraction ϕ for different nanorod stiffness ($K_{\text{stiffness}}$). ($T^* = 1.0$, $\dot{\gamma} = 0.1$).

nanotubes are not completely stiff.^{36,37} The effective aspect ratio of filler gradually increases with its stiffness, which reduces the percolation threshold of PNCs.^{13,27,38–40,44} Under the shear field, it induces the destruction and build-up of the conductive network. CNTs in a shear gradient can pick up other CNTs and stick to each other, which is called the ‘picking-up’ mechanism.¹⁴ Meanwhile, the decrease or the increase of the electrical conductivity depends on the initial dispersion state of filler. For

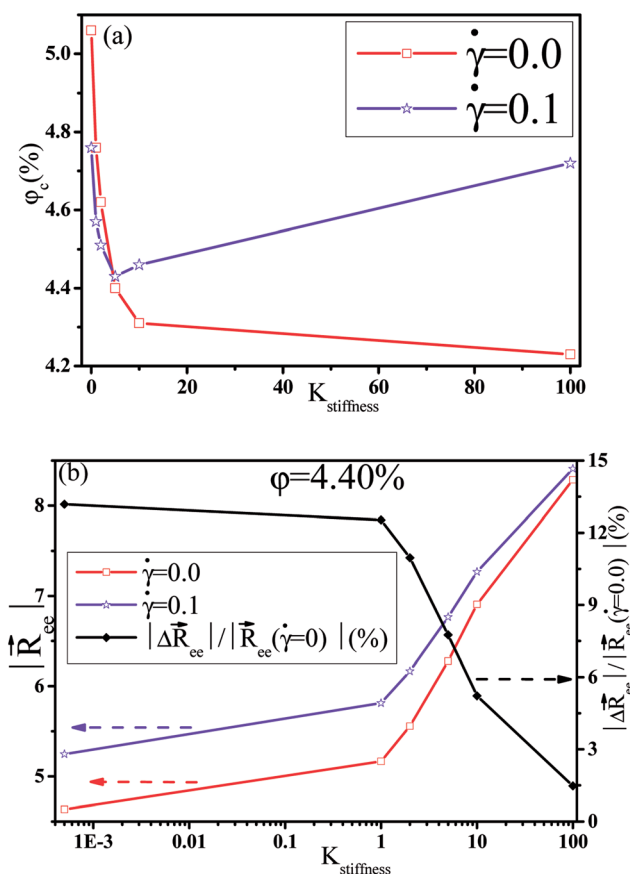


Fig. 7 (a) The percolation threshold ϕ_c for different nanorod stiffness ($K_{\text{stiffness}}$) at shear rates $\dot{\gamma} = 0.0$ and 0.1 . (b) The left axis denotes the end-to-end distance $|\bar{R}_{ee}|$ of the nanorod at shear rates $\dot{\gamma} = 0.0$ and 0.1 , while the right axis represents the change of $|\Delta \bar{R}_{ee}| / |\bar{R}_{ee}(\dot{\gamma} = 0.0)| = |\bar{R}_{ee}(\dot{\gamma} = 0.1) - \bar{R}_{ee}(\dot{\gamma} = 0.0)| / |\bar{R}_{ee}(\dot{\gamma} = 0.0)|$ with different stiffness ($K_{\text{stiffness}}$) ($\phi = 4.40\%$). ($T^* = 1.0$, $\dot{\gamma} = 0.1$).

the well dispersed CNTs in PC matrix, the interconnected conductive network changes the materials from insulator to the conductor, which is attributed to the agglomeration of CNTs under steady shear field.¹⁴ However, for the initially partially aggregated filler, the breakage of the conductive network significantly reduces the electrical conductivity.^{53,63} In total, the shear-induced change of CNT structure (such as CNT alignment^{29,64} and the CNT entanglement⁶⁵) is responsible for the electrical conductivity. In this simulation, we mainly investigated the effect of the NR stiffness on the conductive behavior of PNCs under the shear field. Currently, only some models^{26,28,38–40} reported the effect of NR stiffness on the percolation threshold of PNCs where the NRs changes from very flexibility to rigidity, which is consistent with ours. For the NR in our simulation, the percolation threshold of PNCs varies from 5.06% to 4.23% with the increase of $K_{\text{stiffness}}$ from 0 to 100 where the effective aspect ratio of NR increases from 5.6 to 9.3. The difference of the percolation threshold for different flexible NRs is 0.83%. In experiments,⁶⁶ the percolation threshold decreases from 8.3% to 5.9% when the aspect ratio of silver nanowire increases from 9 to 16 in silver nanowire-polystyrene nanocomposites. This difference of the percolation threshold is larger than ours. On



the one hand, the NR in the simulation is short.^{8,67} The change of the effective aspect ratio of the short NR is limited with different stiffness and different shear rates. On the other hand, in the simulation, we adopted the conductive probability to qualitatively reflect the electrical conductivity in experiments, not quantitatively. Meanwhile, it just changes from 0.0 to 1.0 (one order of magnitude). However, in experiments,² the electronic conductivity is used, which can change over more than 10 orders of magnitude near the percolation threshold. Under the shear field, for the rigid NR with $K_{\text{stiffness}} = 100$, the percolation threshold increases from 4.23% at $\dot{\gamma} = 0.0$ to 4.72% at $\dot{\gamma} = 0.1$. However, similarly the difference is also small compared with that in experiment.¹³ Besides the above two possibilities, the NR dispersion is relatively uniform in the simulation, which forms a loose conductive network. In experiments,¹³ the closely connected conductive network is formed. Under the shear field, accompanied by the destruction of the closely connected conductive network, the significant decrease of the electronic conductivity leads to the large increase of the percolation threshold. However, these are three possible reasons for small difference of the percolation threshold in our systems, which are expected to be proved by using long NRs in the future. In total, compared with the previous work, the main new phenomena revealed here are summarized below: (1) in the quiescent state, the conductive probability of PNCs gradually increases with the increase of the NR stiffness; however, under the shear field, it first increases and then decreases. (2) Compared with in the quiescent state, the decrease or the increase of the conductive probability under the shear field depends on the NR stiffness. (3) For flexible NRs, the conductive probability first increases and then decreases with the increase of the shear rate. As a result, the conductive probability reaches maximum at the mediate shear rate, which is consistent with the maximum effective aspect ratio of NRs. These results can help to further understand the effect of NR stiffness on the conductive property of PNCs under the shear field.

4. Conclusions

In this work, the effect of nanorod (NR) stiffness on the conductive probability of NR filled polymer nanocomposites (PNCs) has been investigated under the quiescent state and the shear field. In the quiescent state, the conductive probability of PNCs gradually increases with increasing the NR stiffness, which is reflected by the low percolation threshold; however, under the shear field, it first increases and then decreases. The largest conductive probability appears at the moderate NR stiffness, which result from the competition between the improved effective aspect ratio of NR and the breakage of conductive network induced by the shear field. In addition, compared with in the quiescent state, the NR stiffness determines the decrease or the increase of the conductive probability under the shear field. At low NR stiffness, the improved effective aspect ratio of NR enhances the conductive probability. While at high NR stiffness, the conductive network is significantly broken down, which reduces the conductive probability. At low shear rate, NRs with low stiffness first straighten, which

increases the effective aspect ratio; however, at high shear rate, the increase of effective aspect ratio is limited. As a result, the maximum effective aspect ratio of NRs appears at the moderate shear rate, leading to the maximum conductive probability at the mediate shear rate. In summary, this work can further help to understand the effect of NR stiffness on the conductive property of PNCs under the shear field.

Conflicts of interest

There are no conflicts to declare.

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References

- 1 B. Guo, Z. Tang and L. Zhang, *Prog. Polym. Sci.*, 2016, **61**, 29–66.
- 2 R. M. Mutiso and K. I. Winey, *Prog. Polym. Sci.*, 2015, **40**, 63–84.
- 3 R. Haggemueller, C. Guthy, J. R. Lukes, J. E. Fischer and K. I. Winey, *Macromolecules*, 2007, **40**, 2417–2421.
- 4 M. Foygel, R. D. Morris, D. Anez, S. French and V. L. Sobolev, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2005, **71**, 104201.
- 5 Y. Shudo, A. Izumi, K. Hagita, T. Nakao and M. Shibayama, *Polymer*, 2017, **116**, 506–514.
- 6 I. Balberg, *Philos. Mag. B*, 1987, **56**, 991–1003.
- 7 I. Alig, P. Pötschke, D. Lellinger, T. Skipa, S. Pegel, G. R. Kasaliwal and T. Villmow, *Polymer*, 2012, **53**, 4–28.
- 8 W. Bauhofer and J. Z. Kovacs, *Compos. Sci. Technol.*, 2009, **69**, 1486–1498.
- 9 Y. Zeng, P. Liu, J. Du, L. Zhao, P. M. Ajayan and H.-M. Cheng, *Carbon*, 2010, **48**, 3551–3558.
- 10 Y. Bréchet, J. Y. Cavallé, E. Chabert, L. Chazeau, R. Dendievel, L. Flandin and C. Gauthier, *Adv. Eng. Mater.*, 2001, **3**, 571–577.
- 11 J. Li, P. C. Ma, W. S. Chow, C. K. To, B. Z. Tang and J. K. Kim, *Adv. Funct. Mater.*, 2007, **17**, 3207–3215.
- 12 S. B. Kharchenko, J. F. Douglas, J. Obrzut, E. A. Grulke and K. B. Migler, *Nat. Mater.*, 2004, **3**, 564–568.
- 13 J. Obrzut, J. F. Douglas, S. B. Kharchenko and K. B. Migler, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2007, **76**, 195420.
- 14 T. Skipa, D. Lellinger, W. Böhm, M. Saphiannikova and I. Alig, *Polymer*, 2010, **51**, 201–210.



- 15 S. Kale, F. A. Sabet, I. Jasiuk and M. Ostoj-Starzewski, *J. Appl. Phys.*, 2016, **120**, 045105.
- 16 S. Kale, F. A. Sabet, I. Jasiuk and M. Ostoj-Starzewski, *J. Appl. Phys.*, 2015, **118**, 154306.
- 17 G. Ambrosetti, C. Grimaldi, I. Balberg, T. Maeder, A. Danani and P. Ryser, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2010, **81**, 155434.
- 18 G. Ambrosetti, N. Johnner, C. Grimaldi, T. Maeder, P. Ryser and A. Danani, *J. Appl. Phys.*, 2009, **106**, 016103.
- 19 C. Li and T. W. Chou, *Appl. Phys. Lett.*, 2007, **90**, 491.
- 20 R. H. Otten and d. S. P. Van, *Phys. Rev. Lett.*, 2009, **103**, 225704.
- 21 A. V. Kyrlyuk and P. V. D. Schoot, *Proc. Natl. Acad. Sci. U. S. A.*, 2008, **105**, 8221–8226.
- 22 B. Nigro, C. Grimaldi, M. A. Miller, P. Ryser and T. Schilling, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2013, **88**, 042140.
- 23 H. Woo Cho, S. Nam, S. Lim, D. Kim, H. Kim and B. June Sung, *J. Appl. Phys.*, 2014, **115**, 154307.
- 24 H. M. Ma and X. L. Gao, *Polymer*, 2008, **49**, 4230–4238.
- 25 R. M. Mutiso, M. C. Sherrott, J. Li and K. I. Winey, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2013, **86**, 278–281.
- 26 W. S. Bao, S. A. Meguid, Z. H. Zhu, Y. Pan and G. J. Weng, *J. Appl. Phys.*, 2013, **113**, 234313.
- 27 H. M. Ma, X.-L. Gao and T. B. Tolle, *Appl. Phys. Lett.*, 2010, **96**, 061910.
- 28 S. Kwon, H. W. Cho, G. Gwon, H. Kim and B. J. Sung, *Phys. Rev. E*, 2016, **93**, 032501.
- 29 S. Gong, Z. H. Zhu and S. A. Meguid, *Polymer*, 2015, **56**, 498–506.
- 30 A. E. Eken, E. J. Tozzi, D. J. Klingenberg and W. Bauhofer, *J. Appl. Phys.*, 2011, **109**, 084342.
- 31 C. Jizhe and N. Mohammad, *Modell. Simul. Mater. Sci. Eng.*, 2016, **24**, 065004.
- 32 F. Du, J. E. Fischer and K. I. Winey, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2005, **72**, 121404.
- 33 G. Kwon, Y. Heo, K. Shin and B. J. Sung, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2012, **85**, 011143.
- 34 A. E. Eken, E. J. Tozzi, D. J. Klingenberg and W. Bauhofer, *Polymer*, 2012, **53**, 4493–4500.
- 35 A. E. Eken, E. J. Tozzi, D. J. Klingenberg and W. Bauhofer, *Polymer*, 2011, **52**, 5178–5185.
- 36 F. Dalmas, R. Dendievel, L. Chazeau, J.-Y. Cavallé and C. Gauthier, *Acta Mater.*, 2006, **54**, 2923–2931.
- 37 Y. B. Yi, L. Berhan and A. M. Sastry, *J. Appl. Phys.*, 2004, **96**, 1318–1327.
- 38 W. Lu, T.-W. Chou and E. T. Thostenson, *Appl. Phys. Lett.*, 2010, **96**, 223106.
- 39 C. Li, E. T. Thostenson and T.-W. Chou, *Compos. Sci. Technol.*, 2008, **68**, 1445–1452.
- 40 L. Berhan and A. M. Sastry, *Phys. Rev. E: Stat., Nonlinear, Soft Matter Phys.*, 2007, **75**, 041121.
- 41 K. Kremer and G. S. Grest, *J. Chem. Phys.*, 1990, **92**, 5057–5086.
- 42 G. N. Toepperwein, N. C. Karayiannis, R. A. Riggleman, M. Kröger and J. J. D. Pablo, *Macromolecules*, 2011, **44**, 1034–1045.
- 43 G. N. Toepperwein, R. A. Riggleman and J. J. D. Pablo, *Macromolecules*, 2012, **45**, 543–554.
- 44 Y. Gao, D. Cao, Y. Wu, J. Liu and L. Zhang, *Polymer*, 2016, **101**, 395–405.
- 45 Y. Gao, D. Cao, J. Liu, J. Shen, Y. Wu and L. Zhang, *Phys. Chem. Chem. Phys.*, 2015, **17**, 22959–22968.
- 46 Y. Gao, J. Liu, J. Shen, L. Zhang, Z. Guo and D. Cao, *Phys. Chem. Chem. Phys.*, 2014, **16**, 16039–16048.
- 47 S. Plimpton, *J. Comput. Phys.*, 1995, **117**, 1–19.
- 48 Y. Wang, G. J. Weng, S. A. Meguid and A. M. Hamouda, *J. Appl. Phys.*, 2014, **115**, 193706.
- 49 C. Li, E. T. Thostenson and T.-W. Chou, *Appl. Phys. Lett.*, 2007, **91**, 223114.
- 50 Y. Feng, N. Ning, Q. Zhao, J. Liu, L. Zhang, M. Tian and J. Mi, *Soft Matter*, 2014, **10**, 8236.
- 51 S. Nam, H. W. Cho, S. Lim, D. Kim, H. Kim and B. J. Sung, *ACS Nano*, 2013, **7**, 851–856.
- 52 I. Balberg, C. H. Anderson, S. Alexander and N. Wagner, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 1984, **30**, 3933–3943.
- 53 I. Alig, D. Lellinger, M. Engel, T. Skipa and P. Pötschke, *Polymer*, 2008, **49**, 1902–1909.
- 54 M. E. Tuckerman, C. J. Mundy, S. Balasubramanian and M. L. Klein, *J. Chem. Phys.*, 1997, **106**, 5615–5621.
- 55 A. W. Lees and S. F. Edwards, *J. Phys. C: Solid State Phys.*, 1972, **5**, 1921.
- 56 P. C. Hiemenz and T. P. Lodge, in *Polymer chemistry*, CRC Press, London, New York, 2007, p. 587.
- 57 R. G. Larson, *The Structure and Rheology of Complex Fluids*, Oxford University Press, New York, 1999, p. 663.
- 58 A. L. Frischknecht, M. J. A. Hore, J. Ford and R. J. Composto, *Macromolecules*, 2013, **46**, 2856–2869.
- 59 A. Rungta, B. Natarajan, T. Neely, D. Dukes, L. S. Schadler and B. C. Benicewicz, *Macromolecules*, 2012, **45**, 9303–9311.
- 60 S. K. Kumar, N. Jouault, B. Benicewicz and T. Neely, *Macromolecules*, 2013, **46**, 3199–3214.
- 61 J. F. Moll, P. Akcora, A. Rungta, S. Gong, R. H. Colby, B. C. Benicewicz and S. K. Kumar, *Macromolecules*, 2011, **44**, 7473–7477.
- 62 V. Kumar and A. Rawal, *Polymer*, 2016, **97**, 295–299.
- 63 I. Alig, T. Skipa, M. Engel, D. Lellinger, S. Pegel and P. Pötschke, *Phys. Status Solidi B*, 2007, **244**, 4223–4226.
- 64 J. Xu, W. Florkowski, R. Gerhardt, K.-s. Moon and C.-P. Wong, *J. Phys. Chem. B*, 2006, **110**, 12289–12292.
- 65 R. H. Schmidt, I. A. Kinloch, A. N. Burgess and A. H. Windle, *Langmuir*, 2007, **23**, 5707–5712.
- 66 S. I. White, R. M. Mutiso, P. M. Vora, D. Jahnke, S. Hsu, J. M. Kikkawa, J. Li, J. E. Fischer and K. I. Winey, *Adv. Funct. Mater.*, 2010, **20**, 2709–2716.
- 67 D. A. T. Nikta Fakhri, L. Cognet, R. Bruce Weisman and M. Pasquali, *Proc. Natl. Acad. Sci. U. S. A.*, 2009, **106**, 14219–14223.

