





Cite this: *RSC Adv.*, 2017, 7, 54734

Tunable thermal conductivities of graphene and graphyne under in-plane torsion†

Yinfeng Li, *^{ab} Qianling Lin,^a Ye Li^b and Daxiang Cui ^c

Using the non-equilibrium molecular dynamics method, the thermal properties of two dimensional nanomaterials are investigated by considering graphene and graphyne nanosheets with circular boundaries. The thermal transport efficiency of graphene and graphyne under heat flux from the inner boundary to outer boundary is revealed to be tunable by applying in-plane torsion at the inner boundary, and the tunable range of thermal conductivity for graphyne could be up to 37% (15% for graphene). With the increase of rotation angle, the thermal conductivities of both graphene and graphyne are found to increase at small rotation angles and then decrease after the occurrence of wrinkle deformation. The maximum thermal conductivity appears at the onset of wrinkling which depends on the lattice structure and stiffness of the nanosheets. By systematically investigating the morphological characteristics and the phonon spectra under different torsion angles, the tunable thermal conductivities of both graphene and graphyne are found to be controlled by three factors including surface smoothness, stress concentration and lattice instability. The increase of thermal conductivity with small torsion angles is caused by the suppressed surface fluctuation which decreases the phonon scattering, while the wrinkling and lattice instability occurring under large torsion angles accounts for the deterioration of thermal conductivity. Since the fluctuation of graphyne is efficiently compressed at smaller torsion angles compared to graphene, the maximum thermal conductivity of graphyne appears earlier than graphene. Such correlation between out-of-plane deformation and in-plane thermal conductivity provides new insights into the thermal management of two dimensional nanomaterials.

Received 9th October 2017
 Accepted 20th November 2017

DOI: 10.1039/c7ra11128b

rsc.li/rsc-advances

Introduction

Graphene, as an ideal two dimensional (2D) nanomaterial composed of a single layer of atoms organized in a hexagonal lattice, has attracted wide attention since it was fabricated in 2004.¹ Owing to their extraordinary mechanical, thermal, electronic and optical characteristics,^{2–4} researchers are actively exploiting 2D materials for various applications in next generation electronics,^{5,6} microchips,⁷ composites,^{8,9} biosensors, drug delivery,¹⁰ and energy harvesting and conversion systems.¹¹ Graphyne is another form of carbon nanosheet composed of sp and sp² carbon atoms, which was first predicted by Baughman *et al.*¹² New synthetic methods enable the successful fabrication of graphyne on copper substrates *via* a cross-coupling reaction using hexaethynylbenzene.^{13–15}

Nowadays, graphyne has become the focus of new investigations because of its promising applications in novel nano-electronic and optoelectronic devices with highly tunable band gaps, large carrier mobility, and unique transport properties.^{16–19}

Owing to the structural features, 2D nanomaterials are easily susceptible to out-of-plane deformation and tend to form wrinkles and corrugations during fabrication and application.^{20–22} These out-of-plane deformations can be seen as a stability mechanism to relax the in-plane strain of thin 2D materials,^{23,24} and have been reported to affect the electronic and chemical properties by inducing effective magnetic fields and changing local potentials.^{25–27} Unusual spiral wrinkling patterns has been reported for graphene nanosheets under circular shearing at the inner edge,^{28,29} and show potential applications as nanosensor, tunable magnetic or electronic devices, as well as patterned stretchable electronics.^{30–32} The wrinkle characteristic and torque capacity of circular graphene and graphyne under in-plane torsion has been studied by varying approaches. Zhang *et al.* studied the formation and characteristics of the wrinkles in annular graphene under in-plane rotation combing grillage model and molecular dynamics simulation.^{32,33} Qin *et al.* investigated the effect of wrinkles on the surface area of graphene using classical

^aDepartment of Engineering Mechanics, School of Naval Architecture, Ocean and Civil Engineering, State Key Laboratory of Ocean Engineering, Shanghai Jiao Tong University, Shanghai 200240, China. E-mail: liyinfeng@sjtu.edu.cn

^bCollaborative Innovation Center for Advanced Ship and Deep-Sea Exploration, Shanghai 200240, China

^cInstitute of Nano Biomedicine and Engineering, Shanghai Jiao Tong University, Shanghai 200240, China

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c7ra11128b



molecular dynamics simulations, and suggested that the high specific surface area of graphene can only be affected up to 2% regardless of loading conditions, geometry, and defects.³⁴ Li *et al.* reported the wrinkling characteristics of surface functionalized graphene and graphyne under circular shearing at the inner edge.^{29,35}

The keep miniaturization of electronic devices raises a strong demand for high efficient heat dissipation and thermal management to ensure device performance. The thermal conductivity of graphene has been revealed to be sensitive to external loadings such as tension and out-of-plane rotation,^{36,37} and the largest enhancement of thermal conductivity provided by applied strain is reported up to 40%.³⁸ The effects of non-uniform strain with specific distribution has also been revealed to be associated with the electronic structure and Raman spectra of graphene.³⁹ Thus, in addition to its mechanical response under circular shearing, thermal conductivity of 2D nanomaterials tuned by in-plane torsion is also of crucial interest for its design and practical application. It is necessary to understand how in-plane rotation influences the thermal conductivity of 2D nanomaterials for the design of micro and nano electro-mechanical system devices. In this paper, thermal characteristics of 2D nanomaterials with circular boundaries under in-plane torsion applied at its inner edge are studied using molecular dynamics method. The thermal conductivities of graphene and graphyne annuli under heat flux from inner boundary to outer boundary are calculated at varying in-plane torsion angle. The radial thermal conductivity of both graphene and graphyne shows interesting dependency on the torsion-angle. Systematic analyses on surface morphology and phonon spectra are carried out for the mechanism of such torsion-dependent thermal conductivity. The reported tunable thermal conductivities of graphene and graphyne under in-plane torsion are expected to stand for a wide range of 2D nanomaterials. Our results provide insights for developing next generation nanodevices with tunable thermal conductivity.

Methods

Fig. 1a shows the atomistic model of circular graphene with outer radius $R_o = 10$ nm under in-plane torsion. Gray atoms inside $R_i = 3$ nm and outside R_o represent the inner boundary and outer boundary, and cyan atoms in between represent the free carbon atoms of graphene. The torsion angle $\Delta\theta$ applied at the inner radius equals to 10° . Adaptive Intermolecular Reactive Empirical Bond Order (AIREBO) potential⁴⁰ implemented in the LAMMPS package⁴¹ is adopted for the carbon-carbon interactions. Prior to rotation, a relaxation process of 300 000 MD steps with time step $\Delta t = 0.1$ fs is carried out for equilibrium. Boundary atoms are free to move in the plane of graphene while the out-of-plane displacement is constrained by enforcing zero force and velocity along out-of-plane direction. During the rotation process, the atoms inside the inner boundary R_i are rotated around the center as a rigid body at constant angular velocity 0.1 rad ps^{-1} with outer boundary fixed. After rotation, another relaxation process of 500 000 MD steps with fixed inner

and outer boundary is enforced. The relaxing and loading process is performed at room temperature (300 K) under NVT ensemble. Thermal conductivity is then computed by non-equilibrium molecular dynamics method by dividing the free atoms of annulus into concentric loops, where the most inner loop is assigned to be the hot region with fixed temperature 350 K while the most outer loop is the cold one with fixed 250 K. Heat flux J can be calculated from the exchanging of kinetic energy between the cold and hot domains by the equation:

$$J = \frac{1}{V} \left[\sum_i \varepsilon_i v_i + \sum_{ij:i \neq j} (F_{ij} v_i) + \frac{1}{6} \sum_{ijk:i \neq j \neq k} (F_{ijk} v_i)(r_{ij} + r_{ik}) \right] \quad (1)$$

where the subscripts i, j, k denote the three different atoms i, j, k . ε_i and v_i are the energy and velocity of atom i . Vector r_{ij} represents the interatomic distance between atoms i and j , and F_{ij} and F_{ijk} represent the two-body and three-body force, respectively. V is the volume of the studied system.

The whole exchanging process is performed under NVE ensemble for 1 ns to reach a steady temperature gradient along radial direction. The temperatures of each loop are determined by:

$$T_{\text{slab}} = \frac{2}{3Nk_B} \sum_j \frac{p_j^2}{2m_j} \quad (2)$$

where N is the number of carbon atoms in this loop, k_B is Boltzmann's constant and p_j is the momentum of atom j . The temperature profile of circular graphene $T(r)$ can be obtained by averaging the temperature of each loop over 50 ps. Fig. 1b displays the temperature profile of a circular graphene as a function of atomic position along the radial direction. For an equilibrium graphene with fixed temperature at its inner and outer boundaries, the heat equation is given as:

$$\Delta T = \frac{1}{r} \frac{\partial T}{\partial r} + \frac{\partial^2 T}{\partial r^2} = 0, \quad \text{with } T|_{r=r_i} = T_1, \quad T|_{r=r_o} = T_2 \quad (3)$$

By solving eqn (3), we can get the analytic solution of temperature profile in annulus:

$$T = a \ln r + b \quad (4)$$

where a and b are integrating coefficients which can be determined by fitting this equation to the simulated temperature profile. The thermal conductivity could be computed by the Fourier law:

$$k = -J \left(\frac{\partial T}{\partial r} \right)^{-1} = -\frac{Jr}{A} \quad (5)$$

where A is the area of cross section along heat flux.

Results and discussion

We first study the dynamic rotation process of the described graphene under in-plane torsion at inner boundary. The graphene sheet is found to remain flat when torsion angle $\Delta\theta$ is small. With the increase of $\Delta\theta$, the graphene nanosheet buckles



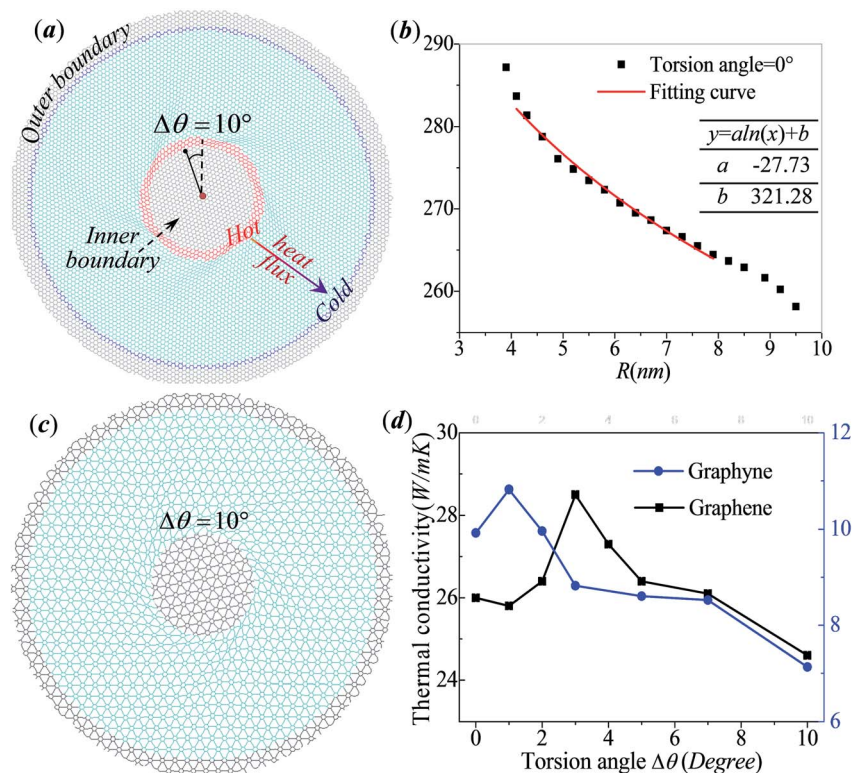


Fig. 1 Thermal conductivities of circular graphene and graphyne under circular in-plane torsion. (a) Atomistic model of graphene annulus with $R_i = 3$ nm and $R_o = 10$ nm under torsion angle $\Delta\theta = 10^\circ$; (b) temperature profile of the graphene plotted in (a) under $\Delta\theta = 0^\circ$ as a function of atomic position along the radial direction; (c) atomistic model of graphyne annulus under torsion angle $\Delta\theta = 10^\circ$; (d) evolution of the radial thermal conductivities of graphene (black curve) and graphyne (blue curve) as a function of torsion angle.

suddenly forming a series of spiral wrinkles surrounding the inner boundary. The radial thermal conductivities of circular graphene under different torsion angles are calculated when heat flux from inner boundary to outer boundary. The black in Fig. 1d shows the evolution of the radial thermal conductivity of graphene as a function of the torsion angle $\Delta\theta$. The thermal conductivity anomalously increases from 26 W mK^{-1} to 28.5 W mK^{-1} with the increase of torsion angle in the range of $0^\circ < \Delta\theta < 3^\circ$ and reach a maximum at $\Delta\theta = 3^\circ$, which is just the critical point that the spiral wrinkling appears. The maximum thermal conductivity is about 10% higher than that of the graphene annulus at rest. After that, the thermal conductivity decreases to 24.6 W mK^{-1} with the increase of torsion angle until the final failure of annulus. The total tunable range is about 15% of the pristine value. Furthermore, we repeat the simulation with smaller temperature difference between hot region (350 K) and cold region (310 K). The calculated evolution of radial thermal conductivity is plotted in the Fig. S1.† The evolution of the graphene thermal conductivity with smaller temperature difference also shows an inversed-V-shaped evolution, which is same with the results of circular graphene under larger temperature gradient. Similar wrinkle characteristics are also observed in graphyne as plotted in Fig. 1c. It is noticeable that our calculated thermal conductivity of graphene under torsion angle $\Delta\theta = 0^\circ$ equals to 26 W mK^{-1} , which is smaller than the experimental values of graphene nanoribbons (GNRs) about

$2500\text{--}5000 \text{ W mK}^{-1}$.^{42,43} Previous simulations have found that the thermal conductivity of the GNRs is sensitive to the size of graphene nanosheet due to phonon scattering.⁴⁴ Since the radial length of circular graphene considered in our approach is much smaller than the ultra-long phonon mean free path of experimentally studied graphene (about $1 \mu\text{m}$), our simulations focuses on the qualitative discussion about the evolution of thermal conductivity with in-plane torsion. The effect of in-plane torsion is demonstrated by the relative difference between thermal conductivity of graphene under different torsion degree instead of the absolute calculated results.

We further study the thermal conductivity of graphyne to see whether the in-plane torsion tuned thermal properties of graphene can be applicable for other two dimensional carbon allotropies with different unit lattice. Fig. 1c shows the atomistic model of circular graphyne under torsion angle $\Delta\theta = 10^\circ$. Graphyne annulus also presents unusual spiral wrinkling patterns around inner edge under circular in-plane rotation and has lower thermal conductivity as reported by Zhang *et al.*⁴⁵ Similar to the evolution of graphene thermal conductivity, the radial thermal conductivity of graphyne also shows an inversed-V-shape evolution. The maximum thermal conductivity occurs at $\Delta\theta = 1^\circ$, which is earlier than graphene whose max maximum value occurs at $\Delta\theta = 3^\circ$. As shown by the blue curve in Fig. 1d, the maximum thermal conductivity (10.8 W mK^{-1}) is about 9% higher than that of the graphyne annulus at rest (9.9 W mK^{-1}).



After that, the thermal conductivity decreases to 7.1 W mK^{-1} with the torsion angle, the maximum tunable range of thermal conductivity is up to 37% when torsion angle increases from 0 to 10° .

To explain the reason why thermal conductivities of both graphene and graphyne increases anomalously under slight torsion while the peak value occurs at different torsion angle for graphene and graphyne, we further analyze the change of surface morphology of graphene and graphyne under torsion. The change of graphene surface smoothness is noticeable even before the occurrence of wrinkling under small torsion angle. Fig. 2 plots the contour map of graphene annulus by plotting the out-of-plane displacement of carbon atoms. The equilibrated graphene under torsion angle $\Delta\theta = 0^\circ$ exhibits a number of random fluctuations. After the graphene undergoes slight in-plane torsion at inner boundary, the surface morphology is smoothed by in-plane strain and the random fluctuations disappear gradually. Such improvement of surface smoothness contributes to the propagation of the phonon in graphene,⁴⁶ and leads to the increase of thermal conductivity. With the increase of torsion angle, the severe in-plane strain plays a more and more important role in softening the phonon modes and increasing the lattice anharmonicity, which causes the reduction of thermal conductivity. Furthermore, the enhanced wrinkling amplitude under large torsion angle induces the lattice instability which also contributes to the reduction of thermal conductivity. The contours of graphyne under different torsion angle are also calculated as shown in Fig. S2.† It can be seen that the graphyne annulus also possesses a number of random fluctuations under $\Delta\theta = 0^\circ$, and the surface morphology is

flattened obviously when $\Delta\theta = 1^\circ$. Due to the difference in lattice structure and stiffness, spiral wrinkling happens earlier in graphyne annulus than that of graphene annulus. Thus, the maximum thermal conductivity appears at a smaller torsion angle than that of graphene. The atomistic structures of circular graphene and graphyne under in-plane torsion angle $\Delta\theta = 0^\circ$ and 3° are plotted in Fig. S3 and S4† for comparison.

We also carried out detailed analyses on the phonon spectra of graphene and graphyne under different $\Delta\theta$ to elucidate the tuning effect of torsion on thermal transport efficiency. The phonon spectrum function $G(\omega)$ can be calculated from the Fourier transform of the velocity autocorrelation function (VACF):

$$G(\omega) = \frac{1}{\sqrt{2\pi}} \int e^{i\omega t} \left\langle \sum_{j=1}^N v_j(t) v_j(0) \right\rangle dt \quad (6)$$

where $v_j(t)$ denotes its velocity at time t , and ω is the vibrational wavenumber. Here, the ensemble average is substituted with time average. Time averaging is performed for the recorded velocity over 50 000 steps after equilibrium.

The Fourier transform of the VACF reveals the amount of energy in vibrations at each frequency, which is proportional to the phonon density of states times the occupation of the modes. As shown in Fig. 3a, the phonon spectra of annular graphene under different torsion angles are calculated. Each phonon spectra owns a main peak (G-band) at frequencies around 50 THz, which is consistent with previous result.⁴⁶ Moreover, the peak frequency shift left with the increase of torsion angle. Fig. 3b shows the phonon spectra of circular graphyne under different torsion angles. The main peak occurs at frequencies around 60 THz.

To present the change of phonon spectra with torsion angle $\Delta\theta$, the main peak frequency and the corresponding amplitude of each spectrum of graphene and graphyne are recorded as shown in Fig. 4. The peak frequencies of both graphene and graphyne decrease with the increase of torsion angle. The change of peak frequency is slight at small torsion angle while the peak frequency drops dramatically under large in-plane torsion. Previous study has reported that the applied strain could soften the G-band (frequency peaks around 50 THz) of the phonon spectra remarkably, causing a reduction in the phonon group velocities.⁴⁶ Hence, the thermal conductivity is reduced according to the classical lattice thermal transport theory:

$$k = \sum_m C_v v_m l \quad (7)$$

where m is the phonon mode occupied at a specific temperature; C_v , v_m and l are the specific heat, group velocity and mean free path of the phonon, respectively. Therefore, the lower decline rate of peak frequency under small torsion angles is reasonable because the relaxed graphene annulus changes from loose to tight during this process.

The black line in Fig. 4b demonstrates that the amplitude of peak for graphene increases with torsion angle when $0^\circ < \Delta\theta < 3^\circ$ while decreases with torsion angle when $\Delta\theta > 3^\circ$. The evolution of amplitude with torsion angle is consistent with that

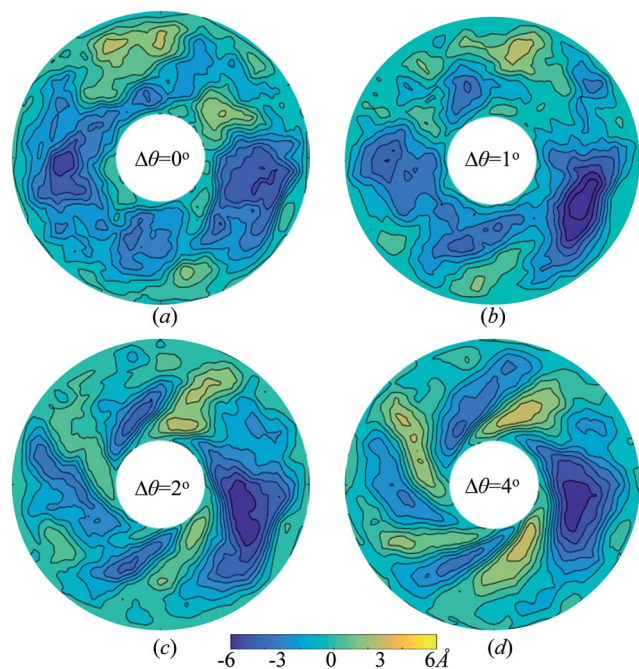


Fig. 2 Change of surface morphology with the increase of torsion angle for circular graphene with $R_i = 3 \text{ nm}$ and $R_o = 10 \text{ nm}$. (a–d) Contour maps of circular graphene annulus under torsion angle $\Delta\theta = 0^\circ, 1^\circ, 2^\circ$ and 3° .



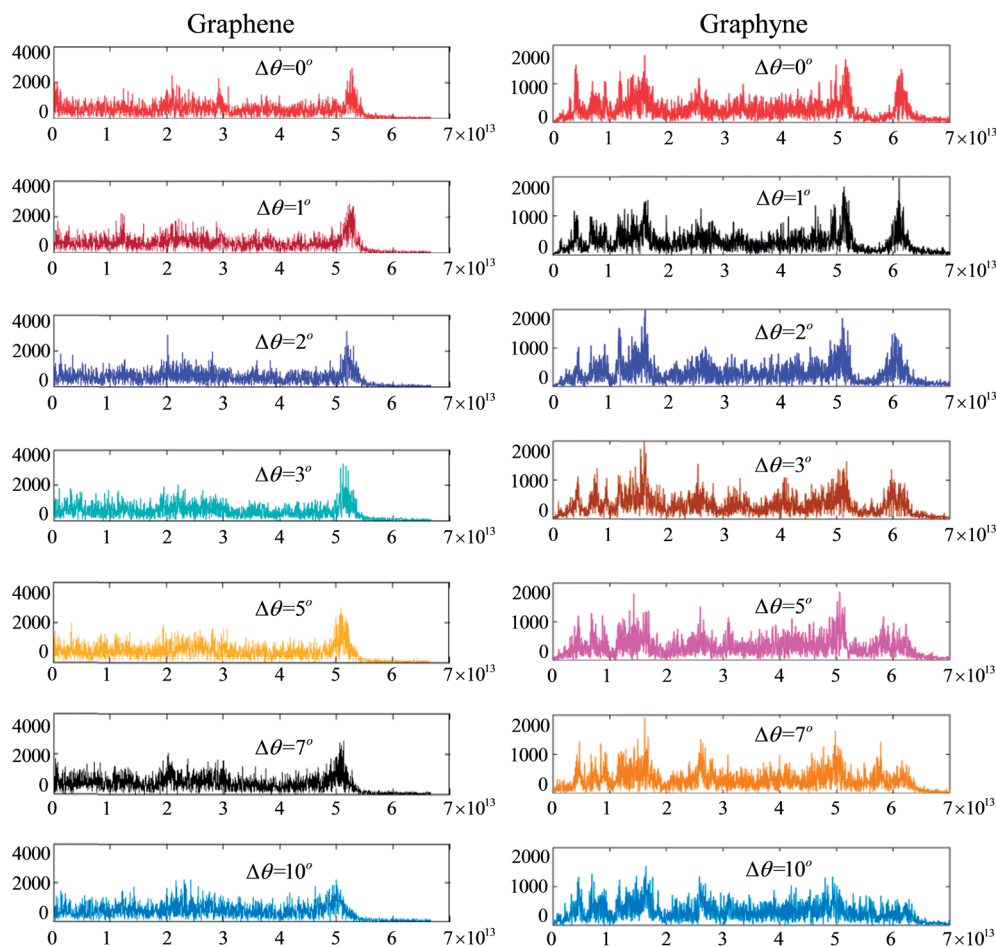


Fig. 3 Phonon spectrum of graphene and graphyne under varying in-plane torsion angle $\Delta\theta$ ranging from 0° to 10° . The thermal conductivity is related with the peak frequency of the phonon spectra as well as the amplitude of peak frequency.

of thermal conductivity with $\Delta\theta$. The variation of the amplitude of peak frequency can be attributed to the change of phonon scattering which has already been suggested as the main factor for the reduction of thermal conductivity. More severe phonon scattering corresponds to lower frequency amplitude. Under small torsion angle, the thermal conductivity increases with the improvement of surface smoothness of graphene annulus, which reduces the degree of phonon scattering. Meanwhile, the strain induced redshift of the peak frequency of the phonon

spectra is too small to change the thermal conductivity, *i.e.*, the decline rate of thermal conductivity due to phonon peak frequency redshift at this stage is small. As a result, the thermal conductivity increases with torsion angle under small torsion angle. With the increasing of torsion angle, the increased in-plane shear strain and lattice instability result in the redshift of peak frequency and the enhancement of phonon scattering. The combined influences provided by the changes of peak frequency and phonon scattering lead to the decrease of

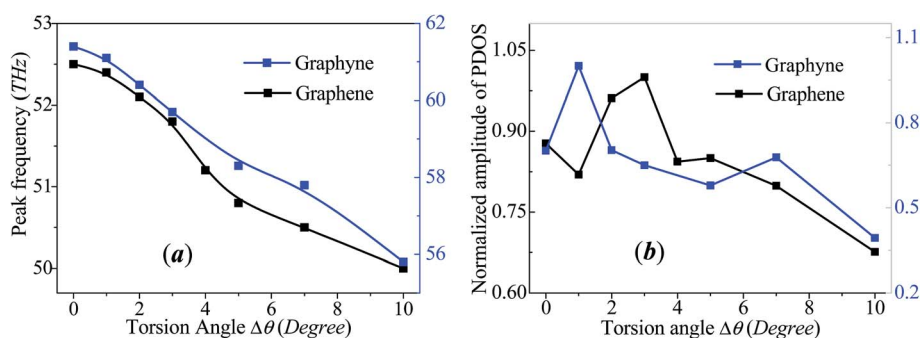


Fig. 4 Evolutions of the peak frequency of the phonon spectra (a) and the amplitude of the peak frequency (b) as a function of the torsion angle for circular graphene and graphyne.



thermal conductivity. The overall change of thermal conductivity follows an inverted V-shaped evolution with the increase of rotation angle. For graphyne represented by the blue line in Fig. 4b, the peak frequency amplitude also shows an inverted V-shaped evolution with torsion angle while the maximum value appears at $\Delta\theta = 1^\circ$. The specified mechanism for the in-plane tuned thermal conductivity is expected to be applicable for 2D nanomaterials of different lattice structures.

Conclusions

In conclusion, the in-plane torsion tuned thermal conductivities of circular graphene and graphyne have been studied systematically. The thermal conductivity shows an interesting inverted V-shaped evolution with the torsion angle, and the tunable range of thermal conductivity for graphyne could be up to 37% (15% for graphene). The evolution of thermal conductivity with torsion angle is interpreted from the perspectives of out-of-plane deformation and the phonon spectra of nanosheets. Under small in-plane torsion angle, the surface morphology of the 2D nanomaterials is flattened and phonon scattering is depressed, which contribute to the increase of thermal conductivity. With the increase of torsion angle, the enhanced in-plane strain causes the occurrence of wrinkling and lattice instability which soften the phonon modes and increases phonon scattering, leading to the deterioration of thermal conductivity. The results show that the in-plane thermal conductivity can be effectively regulated by out-of-plane deformations which affect the phonon spectra. Similar tunable thermal conductivity has also been revealed for graphyne under in-plane torsion. Our results indicate that the described tunable thermal properties as well as the specified mechanism stand for wide range of 2D nanomaterials with different unit lattice structures. The revealed phenomenon and mechanism provides a new route for tailoring the thermal properties of 2D nanosheets, and can also be combined with other approaches, such as surface functionalization, introducing defects and out-of-plane strains for more efficient manipulation of thermal conductivity.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

We gratefully acknowledge the support of the National Natural Science Foundation of China (No. 11402145), National Key Basic Research Program (973 Project) (Project No. 2017YFA0205300). The computational support for this work was provided by Center for HPC, Shanghai Jiao Tong University.

References

- 1 K. S. Novoselov, A. K. Geim, S. V. Morozov, D. Jiang, Y. Zhang, S. V. Dubonos, I. V. Grigorieva and A. A. Firsov, *Science*, 2004, **306**, 666–669.

- 2 L. A. Falkovsky, *J. Phys.: Conf. Ser.*, 2008, **129**, 012004.
- 3 Y. Li, D. Datta and Z. Li, *Carbon*, 2015, **90**, 234–241.
- 4 M. Terrones, A. R. Botello-Méndez, J. Campos-Delgado, F. López-Urías, Y. I. Vega-Cantú, F. J. Rodríguez-Macías, A. L. Elías, E. Muñoz-Sandoval, A. G. Cano-Márquez and J.-C. Charlier, *Nano Today*, 2010, **5**, 351–372.
- 5 D. Boukhvalov, M. Katsnelson and A. Lichtenstein, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2008, **77**, 035427.
- 6 Y. Zhang, Y. W. Tan, H. L. Stormer and P. Kim, *Nature*, 2005, **438**, 201–204.
- 7 S. Cho and M. S. Fuhrer, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2008, **77**, 081402.
- 8 D. W. Boukhvalov and M. I. Katsnelson, *J. Phys.: Condens. Matter*, 2009, **21**, 344205.
- 9 A. Salehi-Khojin, D. Estrada, K. Y. Lin, K. Ran, R. T. Haasch, J.-M. Zuo, E. Pop and R. I. Masel, *Appl. Phys. Lett.*, 2012, **100**, 033111.
- 10 Y. Li, H. Yuan, A. von dem Bussche, M. Creighton, R. H. Hurt, A. B. Kane and H. Gao, *Proc. Natl. Acad. Sci. U. S. A.*, 2013, **110**, 12295–12300.
- 11 H. Gao, *J. Mech. Phys. Solids*, 2014, **62**, 312–339.
- 12 R. H. Baughman, H. Eckhardt and M. Kertesz, *J. Chem. Phys.*, 1987, **87**, 6687–6699.
- 13 E. L. Spitler, C. A. Johnson and M. M. Haley, *Chem. Rev.*, 2006, **106**, 5344–5386.
- 14 C. D. Stevenson, *Acc. Chem. Res.*, 2007, **40**, 703–711.
- 15 G. Li, Y. Li, H. Liu, Y. Guo, Y. Li and D. Zhu, *Chem. Commun.*, 2010, **46**, 3256–3258.
- 16 J. Koo, H. J. Hwang, B. Huang, H. Lee, H. Lee, M. Park, Y. Kwon, S.-H. Wei and H. Lee, *J. Phys. Chem. C*, 2013, **117**, 11960–11967.
- 17 W. Wu, W. Guo and X. C. Zeng, *Nanoscale*, 2013, **5**, 9264–9276.
- 18 Q. Yue, S. Chang, J. Kang, S. Qin and J. Li, *J. Phys. Chem. C*, 2013, **117**, 14804–14811.
- 19 J. Chen, J. Xi, D. Wang and Z. Shuai, *J. Phys. Chem. Lett.*, 2013, **4**, 1443–1448.
- 20 W. Bao, F. Miao, Z. Chen, H. Zhang, W. Jang, C. Dames and C. N. Lau, *Nature Nanotechnol.*, 2009, **4**, 562–566.
- 21 Q. Lu and R. Huang, *Int. J. Appl. Mech. Eng.*, 2009, **1**, 443–467.
- 22 A. Srivastava, C. Galande, L. Ci, L. Song, C. Rai, D. Jariwala, K. F. Kelly and P. M. Ajayan, *Chem. Mater.*, 2010, **22**, 3457–3461.
- 23 J. C. Meyer, A. K. Geim, M. I. Katsnelson, K. S. Novoselov, T. J. Booth and S. Roth, *Nature*, 2007, **446**, 60–63.
- 24 S. Scharfenberg, N. Mansukhani, C. Chialvo, R. L. Weaver and N. Mason, *Appl. Phys. Lett.*, 2012, **100**, 021910.
- 25 K. Eun-Ah and A. H. C. Neto, *EPL*, 2008, **84**, 57007.
- 26 F. Guinea, B. Horovitz and P. Le Doussal, *Phys. Rev. B: Condens. Matter Mater. Phys.*, 2008, **77**, 205421.
- 27 F. Guinea, M. I. Katsnelson and M. A. H. Vozmediano, *Phys. Rev. B: Condens. Matter*, 2008, **77**, 075422.
- 28 Y. Li, Q. Lin and D. Cui, *Sci. Rep.*, 2017, **7**, 41767.
- 29 Y. Li, S. Liu, D. Datta and Z. Li, *Sci. Rep.*, 2015, **5**, 16556.



- 30 J. S. Bunch, S. S. Verbridge, J. S. Alden, A. M. van der Zande, J. M. Parpia, H. G. Craighead and P. L. McEuen, *Nano Lett.*, 2008, **8**, 2458–2462.
- 31 J. S. Bunch, A. M. van der Zande, S. S. Verbridge, I. W. Frank, D. M. Tanenbaum, J. M. Parpia, H. G. Craighead and P. L. McEuen, *Science*, 2007, **315**, 490–493.
- 32 Z. Zhang, W. H. Duan and C. M. Wang, *Nanoscale*, 2012, **4**, 5077–5081.
- 33 Z. Zhang, W. H. Duan and C. M. Wang, *J. Appl. Phys.*, 2013, **113**, 014902.
- 34 Z. Qin, M. Taylor, M. Hwang, K. Bertoldi and M. J. Buehler, *Nano Lett.*, 2014, **14**, 6520–6525.
- 35 Q. Lin, Y. Li and H. Yao, *Phys. Chem. Chem. Phys.*, 2017, **19**, 20049–20056.
- 36 W. Ning, X. Lanqing, W. Hui-Qiong and Z. Jin-Cheng, *Nanotechnology*, 2011, **22**, 105705.
- 37 C. Si, G. Lu, B.-Y. Cao, X.-D. Wang, Z. Fan and Z.-H. Feng, *J. Appl. Phys.*, 2017, **121**, 205102.
- 38 Y. Kuang, L. Lindsay and B. Huang, *Nano Lett.*, 2015, **15**, 6121–6127.
- 39 C. Si, Z. Sun and F. Liu, *Nanoscale*, 2016, **8**, 3207–3217.
- 40 S. J. Stuart, A. B. Tutein and J. A. Harrison, *J. Chem. Phys.*, 2000, **112**, 6472–6486.
- 41 S. Panigrahi, A. Bhattacharya, S. Banerjee and D. Bhattacharyya, *J. Phys. Chem. C*, 2012, **116**, 4374–4379.
- 42 W. W. Cai, A. L. Moore, Y. W. Zhu, X. S. Li, S. S. Chen, L. Shi and R. S. Ruoff, *Nano Lett.*, 2010, **10**, 1645–1651.
- 43 C. Wang, J. Guo, L. Dong, A. Aiyiti, X. Xu and B. Li, *Sci. Rep.*, 2016, **6**, 25334.
- 44 Z. Guo, D. Zhang and X.-G. Gong, *Appl. Phys. Lett.*, 2009, **95**, 163103.
- 45 Y. Y. Zhang, Q. X. Pei and C. M. Wang, *Comput. Mater. Sci.*, 2012, **65**, 406–410.
- 46 W. Ning, X. Lanqing, W. Hui-Qiong and Z. Jin-Cheng, *Nanotechnology*, 2011, **22**, 105705.

