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Introduction

In order to disentangle rapid technological progresses from excessive carbon emission, energy efficiency and recovery is crucial. Amongst many green energy initiatives such as solar cells and wind energy, waste heat harvesting is an area that is often overlooked.^{1–5} Thermoelectrics, which recycles waste heat to electricity, is a key technology cut out for this purpose. At the materials level, the efficiency of a thermoelectrics depends on $zT = S^2 \sigma T/\kappa$, with *S*, σ , and κ denoting Seebeck coefficient, electrical, and thermal conductivity, respectively. Intuitively, a good thermoelectric material should have high Seebeck coefficient and electrical conductivity, with low thermal conductivity. This will enable it to maintain robust temperature gradient while generating enough useful power. However, these properties have negative interdependencies on

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Improved *zT* in Nb₅Ge₃–GeTe thermoelectric nanocomposite[†]

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Robust electronic transport properties is a crucial in designing high performance thermoelectrics. A key similarity between superconductor and thermoelectric lies in their generally high electrical conductivity, even at above its superconducting temperature. In this work, we design a nanocomposite between Nb₅Ge₃ and GeTe-based thermoelectric to improve its thermoelectric figure of merit *zT*. Phase and microstructural characterization shows distinct Nb₅Ge₃ precipitates embed in Ge_{0.9}Sb_{0.1}Te matrix. In addition, experimental electronic and thermal transport analysis, together with density functional theory calculation were employed to show the synergistic effect of doping Sb and Nb₅Ge₃ nanocomposite approach. 10% Sb doping was found to optimize the electronic properties of the GeTe-based matrix. Further addition of 2 wt% Nb₅Ge₃ nanocomposite to the matrix enhances the phonon scattering, which consequently lowers the lattice thermal conductivity, which results in *zT* of up to 2.0 at 723 K. Such superconductor nanocomposite approach shown in this work can be employed to enhance the properties of other thermoelectric materials.

each other, making it an optimization problem. For instance, *S* and σ are interrelated through *n*, carrier concentration *via*:

$$S = \frac{8\pi^2 \kappa_{\rm B}^2 T}{3eh^2} m_{\rm s}^* \left(\frac{\pi}{3n}\right)^{2/3} \tag{1}$$

$$\sigma = ne\mu = \frac{ne^2\tau}{m_{\rm I}^*}.$$
 (2)

There are a few limited ways to simultaneously enhance *S* and σ . For instance, by engineering band convergence, a high value of m_s^* (density of states effective mass) relative to m_1^* (inertial effective mass) can be obtained, resulting in enhanced power factor ($S^2\sigma$). Besides band convergence, other strategies to achieve higher performance (*zT*) such as nanostructuring, magnetic inclusions, tuning scattering parameters, defect engineering, and dynamic doping are widely used.^{6–19}

In recent years, germanium telluride (GeTe) has emerged as one of the best medium temperature thermoelectric materials, with power conversion efficiency above 10%.²⁰⁻²³ This is primarily due to its Peierls distortion, which renders low thermal conductivity, combined with its robust electronic transport properties and its favourable band structure.²⁴⁻⁴⁰ Nevertheless, GeTe suffers from excessive hole concentration. Therefore, most of the efforts on GeTe have focused on counter doping to optimize carrier concentration with higher valence cation (*i.e.* Bi or Sb).^{20,41-51} In addition, doping or alloying with Ti, Cr, Bi₂Te₃, PbTe, Cu₂Te, AgBiSe₂, and heat treatment have also

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been explored.^{41,52–63} In addition, extremely rapid cooling using liquid nitrogen have also been reported to enhance GeTe performance.⁶⁴ More recently, resonant states and entropy engineering have also been reported to improve performance.^{65–74} However, the temperature at which the peak performance occurs in GeTe typically lies at the range of 600–800 K. This coincides with its rhombohedral to cubic phase transition. Unfortunately, the phase transition usually results in poor mechanical properties.^{75–77} In addition, the lattice thermal conductivity is generally higher in cubic phase, therefore limiting the maximum performance of cubic GeTe.

Among the numerous strategies to enhance thermoelectric properties, nanocomposite approach has been well studied in other materials system such as BiSbTe, PbTe, and SnTe.78-80 However, such approach has rarely been studied in GeTe to date. This is primarily due to overwhelming focus to either optimize its carrier concentration, or tuning its phase transition temperature. In this work, we report the synergistic effect of nanocomposite approach by adding superconductor Nb₅Ge₃ into Ge_{0.9}Sb_{0.1}Te matrix. Different weight percentage of Nb₅Ge₃ was added and their electronic and thermal transport properties were analyzed. For the matrix phase, 10% Sb was added into GeTe to optimize its carrier concentration.^{45,81} By analyzing its intrinsic electronic transport quality via weighted-mobility, it was evident that due to the high mobility of superconductor even at its normal state, adding Nb₅Ge₃ only slightly deteriorates its weighted-mobility, while greatly reduces its lattice thermal conductivity. This results in overall enhancement in the *zT* of $\text{Ge}_{0.9}\text{Sb}_{0.1}$ Te from 1.7 to 2.0 at 723 K. It is noteworthy that due its high electrical conductivity, the addition of Nb5Ge3 results in minimal decrease in overall electronic transport properties of the nanocomposite. In addition, it also helps to preserve slight excess in cation (Ge), which has been shown to be effective in optimizing hole concentration in



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moelectric materials and devices including machine learning augmented thermoelectrics, 3D-printing thermoelectrics, and electronic waste recycling for thermoelectrics. He also works on thermoelectrics for various applications in wearables, IoT, and aerospace. GeTe.⁸² This work opens up exciting avenue of pairing superconductor and thermoelectric as nanocomposite to enhance thermoelectric properties.

Methods

Bulk pellets of Ge_{0.9}Sb_{0.1}Te (denoted GST from hereon) were separately prepared by mixing a stoichiometric amount of high-purity elements (>99.99%) followed by sealing at high vacuum of 10^{-5} mbar in quartz ampoules. Samples were subsequently melted at 1173 K and were occasionally shaken to ensure homogeneity, and were kept at this temperature for 12 hours, followed by quenching in ice water. The final ingots were then pulverized using ball milling (SPEX 8000D, USA) machine. The ball milling were done in SPEX steel-jacketed tungsten carbide jar with tungsten carbide balls, sealed in a Ar atmosphere. The resulting fine powder was subsequently consolidated using spark plasma sintering (SPS) at 673 K for 10 minutes under a pressure of 50 MPa. Extra caution was taken during the cooling down by slow cooling at 15 K min⁻¹ and slowly releasing the pressure to the die to avoid cracking the sample. The size of the graphite die was 12.7 mm in diameter. The sintering was done at 1 Pa vacuum. The final density of the pellets was tested using Archimedes method (~98% theoretical density). Nb₅Ge₃ was prepared from its raw elemental constituents by ball milling in stainless steel jacketed tungsten carbide jar (SPEX 8000D) and milled for 6 hours. Part of the final powder was consolidated using spark plasma sintering (SPS) at 673 K for 10 minutes for further testing. The rest of the powders were mixed into GeTe powder before SPS. The Hall carrier concentration was measured using ECOPIA HMS-5500 with 0.5 T magnetic field.

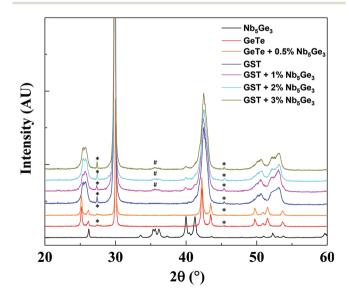


Fig. 1 XRD patterns of pristine GeTe, Nb_5Ge_3 , $Ge_{0.9}Sb_{0.1}Te$ (GST), and GST- Nb_5Ge_3 nanocomposites with different wt%. * denotes Ge precipitates peaks, # denotes Nb_5Ge_3 peaks.

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The Seebeck coefficient and electrical resistivity were measured using ZEM-3 from ULVAC. Thermal conductivity was measured from LFA 457 Netzsch. The heat capacity used for thermal conductivity calculation was estimated using Dulong– Petit approximation. Microscopy analysis was done in field emission scanning electron microscopy FESEM (JEOL JSM 7600F). TEM was collected using Talos F200X. XRD characterization was conducted using D8 Bruker Advance at room temperature. Magnetic property measurements were performed using Superconducting Quantum Interference Device (SQUID, Quantum Design) in a temperature range of 5 to 50 K. The low temperature electrical resistivity was measured using Quantum Design (QD) PPMS (Physical Properties Measurement System) Evercool II TTO (Thermal Transport Option) module.

Band structure calculations were performed using plane-wave self-consistent code implemented in QUANTUM-ESPRESSO.^{83–85} Spin–orbit coupling was taken into consideration during the calculation. The parameterization by Perdew, Burke, and Ernzerhof based on the generalized gradient approximation was used.⁸⁶ Atomic coordinates were fully relaxed until the forces on each

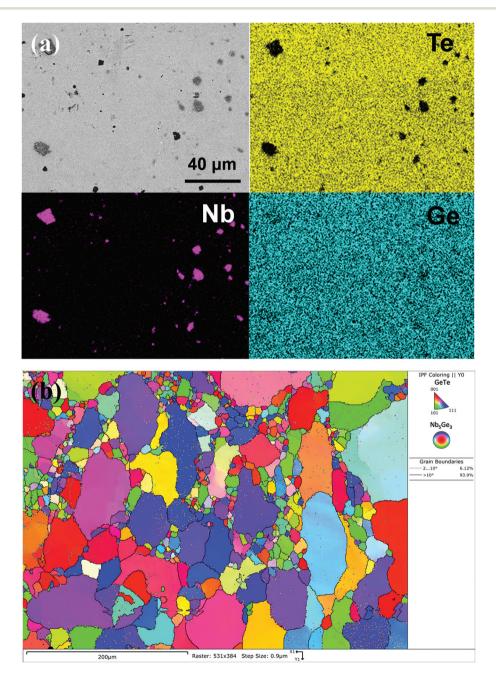


Fig. 2 (a) SEM (scanning electron microscopy) EDS (energy-dispersive X-ray spectroscopy) of the GeTe $-Nb_5Ge_3$ nanocomposite sample showing Nb-rich regions and Te-poor regions, providing clue of Nb_xGe_y phase in GeTe matrix. (b) EBSD (electron backscatter diffraction) mapping showing large grain sizes, with small Nb_5Ge_3 precipitates within the grains.

atom was less than 10–5 eV Å⁻¹. Pristine GeTe in the cubic phase was calculated using the primitive cell. The plane wave cutoff energy was set to 45 Ry. The calculation of GST was based on a $3 \times 3 \times 3$ supercell. The structure was obtained by substituting three Ge atoms with three Sb atom, corresponding to a doping concentration of approximately 11.1%. The plane wave cutoff energy was set to 60 Ry.

Results and discussion

In order to ascertain the phase formation of GST and Nb₅Ge₃, X-ray diffraction (XRD) was done on the final samples, as shown in Fig. 1. It is worth noting that only minute Nb₅Ge₃ peaks can be observed in the GST–Nb₅Ge₃ nanocomposites. This can be attributed to the relatively low volume fraction of Nb₅Ge₃ in GST matrix, considering its much heavier molecular weight (>3 times) compared to GST. In addition, peaks belonging to elemental Ge can be observed at around 28°, consistent with the tendency of GeTe to form Ge precipitates in the matrix.

Further doping of 10% Sb decreases the rhombohedral angle, as evident by the merging of the two peaks around 25° . Importantly, although no Nb₅Ge₃ peak can be observed in the nanocomposite, no shift in peak position or shape can be

observed in the three samples with different fraction of Nb_5Ge_3 , which can be inferred as no Nb_5Ge_3 goes into the lattice of GST.

To further confirm the presence of Nb₅Ge₃ in GST matrix, SEM EDS was done on the GeTe-Nb₅Ge₃ sample, as shown in Fig. 2 and Fig. S5.† Fig. 2(a) shows the presence of dark regions with sizes <10 µm correspond well to Te-poor and Nbrich regions, which provides clues of the Nb_xGe_y phase in the GeTe matrix. It is worth noting that no obvious elemental Ge precipitates can be observed in the figure, which is consistent with the reduced Ge-vacancies as will be elaborated in subsequent figures. In addition to elemental mapping, the EBSD (electron backscatter diffraction) mapping in Fig. 2(b) shows large grain size with area-weighted average size of 46 µm (Fig. S1[†]). In addition, majority of the grains show high-angle boundaries, which has been reported to be effective in reducing thermal conductivity.87 Furthermore, the presence of Nb_xGe_y is further evident in the uniformly distributed precipitates within each grain.

In order to ascertain that the Nb_xGe_y is indeed Nb₅Ge₃ phase in the precipitates, TEM and HRTEM was carried out, as shown in Fig. 3 and Fig. S6.† Fig. 3(a) shows the elemental mapping in TEM, with zoomed in version of Nb₅Ge₃ phase shown in Fig. 3(b). HRTEM image in Fig. 3(c) shows the (100) and (211) planes of Nb₅Ge₃ phase with lattice spacing

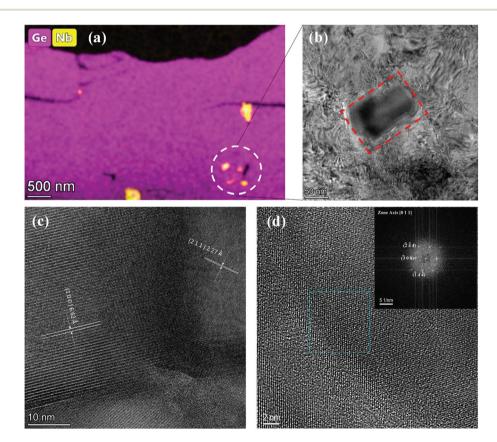


Fig. 3 (a) TEM (transmission electron microscopy) EDS image showing Nb₅Ge₃ phase embed in GeTe matrix. (b) Zoom in image showing Nb₅Ge₃. (c) HRTEM image showing (100) and (211) lattice spacing of Nb₅Ge₃. (d) HRTEM with SAED (selected area electron diffraction) in inset.

of 6.62 Å and 2.27 Å, respectively. The SAED with zone axis (011) is shown in Fig. 3(d), further confirming Nb_5Ge_3 phase signature.

The superconducting properties of Nb₅Ge₃ was characterized using SQUID (superconducting quantum interference device), as shown in Fig. S7.† Temperature dependence of sample magnetization was measured both in zero-field-cooling (ZFC) and field-cooling (FC) processes. In FC process, external magnetic field of 100 Oe was used. The diamagnetic Meissner signal was clear below the temperature ~16 K, indicating the onset of superconducting transition. Magnetic hysteresis loop was measured at 5 K. Since a superconductor is a perfect diamagnet, the sign of magnetic moment is always opposite to that of the external field. Both the Meisnner state and the mixed state can be observed in the loop, exhibiting typical hysteresis loop of type II superconductor. The $T_{\rm C}$ of 16 K in this work is consistent with previous reports on Nb₅Ge₃ with interstitial carbon doping.⁸⁸

With the Nb₅Ge₃ precipitate phase confirmed by both electron microscopy as well as electronic transport, its effect on thermoelectric properties of GST matrix is subsequently examined, as shown in Fig. 4. Evidently, adding Nb₅Ge₃ into pristine GeTe matrix results in slight increase in Seebeck coefficient, as shown in Fig. 4(a). This is consistent with lower carrier concentration in the sample due to less Ge-vacancies (corresponds to less Ge-precipitates), as confirmed by EDS mapping in Fig. 2 which shows the absence of Ge-precipitates. Further doping of 10% Sb increases the Seebeck coefficient of the samples, with negligible difference between samples with different $\mathrm{Nb}_5\mathrm{Ge}_3$ amount.

Fig. 4(b) shows a general trend of increasing resistivity across the entire temperature range with increasing Nb₅Ge₃ wt%, which is also consistent with power factor trend shown in Fig. 4(c). In terms of intrinsic electronic properties, the weighted-mobility, which takes into account the negative interdependencies between Seebeck coefficient and electrical resistivity, is shown in Fig. 4(d). It is evident that while pristine GeTe matrix shows clear acoustic phonon dominated transport, samples with 10% Sb doping show a weaker trend. In addition, introduction of Nb₅Ge₃ only result in small reduction in weighted mobility of all samples, which can be partly attributed to its highly conductive nature. Furthermore, the peaks in weighted mobility at around 650 K can be attributed to rhombohedral-cubic structural phase transition, which results in the convergence of Σ and L band, increasing the overall Seebeck coefficients.

Fig. 5(a) and (b) shows the effect of Sb doping in reducing the hole concentration in GeTe. It is worth noting that the addition of Sb to pristine GeTe results in lowered conduction band, which consequently lead to lowered band gap (from 0.32 eV to 0.15 eV). This is also consistent with the trend of Seeebeck coefficient shown in Fig. 4(a) which shows the downturn of Seebeck coefficients for Sb-doped samples above 723 K. The decrease in Seebeck coefficient at higher temperature can be attributed to thermally activated electron-hole pairs, which results in bipolar conduction, diminishing the net Seebeck coefficient. In contrast, the Seebeck coefficients for pristine GeTe matrix remains in uptrend up to the maximum measure-

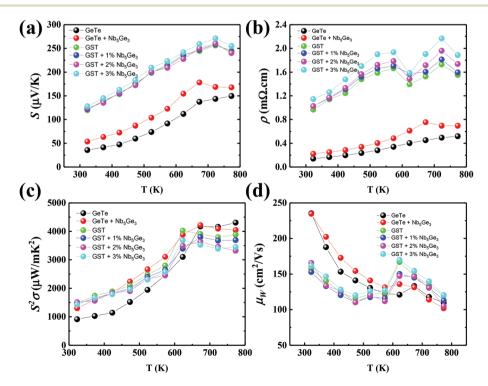


Fig. 4 Temperature dependent (a) Seebeck coefficient (b) electrical resistivity (c) power factor, and (d) weighted-mobility of all samples.

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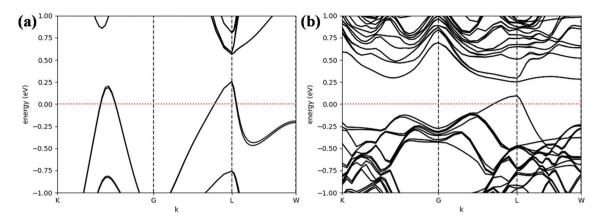


Fig. 5 (a) Electronic band structure of cubic GeTe and (b) cubic Ge_{0.9}Sb_{0.1}Te.

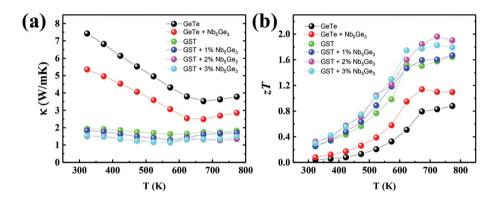


Fig. 6 Temperature dependent (a) total thermal conductivity of all samples. (b) Figure of merit zT of all samples.

ment temperature, which can be associated with its higher band gap.

In terms of thermal transport, the addition of Sb and Nb₅Ge₃ greatly reduces the thermal conductivity, as evident in Fig. 6(a), which subsequently result in enhanced zT of up to 2.0 at 723 K, as shown in Fig. 6(b). To probe deeper into the origin of the thermal conductivity reduction, it is useful to look into the electronic and lattice contribution to thermal conductivity in all samples. Using Seebeck coefficients data together with electrical resistivity of all samples, the Lorenz number and the electronic contribution to thermal conductivity can be estimated (Fig. S2 and S3[†]). Furthermore, the lattice thermal conductivity can then be determined by subtracting the electronic thermal conductivity from the total thermal conductivity, as shown in Fig. 7(a). Evidently, the addition of Nb₅Ge₃ systematically reduces the lattice thermal conductivity, which can be used as an independent tuning knob to enhance *zT*.

While it is arguable that addition of Nb_5Ge_3 also decreases the weighted-mobility of the samples, the net effect of which can be examined by plotting it against the lattice thermal conductivity, as illustrated in Fig. 7(b).

In samples with pristine GeTe matrix, the addition of Nb_5Ge_3 results in systematic downshift of lattice thermal con-

ductivity, as shown by the black arrow. In contrast, samples with $Ge_{0.9}Sb_{0.1}Te$ matrix show consistently low lattice thermal conductivity across a range of weighted-mobility, which lends strong support to the role of point defects introduced by Sb dopants in scattering phonons. More importantly, the addition of Nb₅Ge₃ nanocomposite further reduces the lattice thermal conductivity (from green region to blue region) for the same range of weighted-mobility in Nb₅Ge₃-Ge_{0.9}Sb_{0.1}Te nanocomposites.

Finally, the role of grain boundaries, point defects, and nanoprecipitates in reducing lattice thermal conductivity is laid out in Fig. 7(c). It is evident that due to the large grain sizes (46 μ m), the grain boundaries play negligible effect in reducing lattice thermal conductivity, consistent with the phonon relaxation time in Fig. S4.† On the other hand, by taking into account point defects due to 10% Sb, the spectral thermal conductivity in Fig. 7(c) is drastically reduced, especially in the high frequency range, resulting in reduced lattice thermal conductivity in Fig. 7(d) (dotted blue curve). Last but not least, nanoprecipitates of Nb₅Ge₃ further reduces the spectral thermal conductivity throughout a wide frequency range, resulting in further reduced lattice thermal conductivity (dotted red curve in Fig. 7(d)).

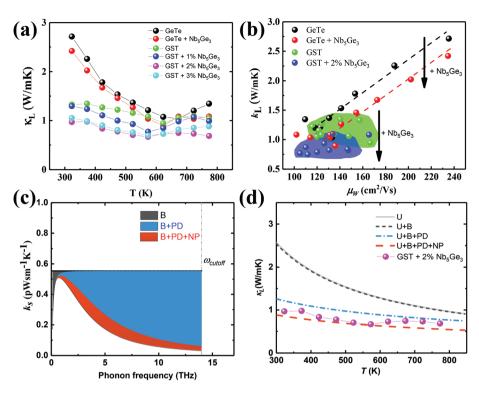


Fig. 7 (a) Temperature dependent lattice thermal conductivity of all samples. (b) Lattice thermal conductivity vs. weighted-mobility relation. (c) Spectral thermal conductivity as a function of phonon frequency based on simple Debye–Callaway model taking into account grain boundaries (B), point defects (B + PD), and nanoprecipitates (B + PD + NP) scattering. (d) Modelled lattice thermal conductivity in comparison to experimental values.

Conclusions

In this work, we demonstrate a novel strategy of using nanocomposite approach to enhance zT in thermoelectric matrix. Nb₅Ge₃ precipitates was added to GeTe-based matrix to form the nanocomposites. Further addition of 10% Sb into the GeTe matrix results in optimized doping and hence power factor. Interestingly, Further analysis using simplified Debye– Callaway model and lattice thermal conductivity *versus* weighted-mobility trend clearly demonstrate the benefits of Nb₅Ge₃ in reducing κ_L far outweigh the reduction in weightedmobility, resulting in enhanced overall zT from 1.7 to 2.0 at 723 K of the nanocomposite. The strategy demonstrated in this work can be used to enhance not only other thermoelectric materials, but also electronic properties of other functional materials in general.

Conflicts of interest

The authors declare no competing financial interest.

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