

Thermomechanical and Thermal Contact Characteristics of Bismuth Telluride Films Electrodeposited on Carbon Nanotube Arrays

By Himanshu Mishra, Baratunde A. Cola, Vijay Rawat, Placidus B. Amama, Kalapi G. Biswas, Xianfan Xu, Timothy S. Fisher, and Timothy D. Sands*

A miniaturized thermoelectric (TE) cooler module is composed of a large number of TE legs connected electrically in series and thermally in parallel.^[1] TE devices operating near room temperature typically create a temperature difference of 30–50 °C, and the TE film thickness for such devices ranges from 10 to 100 μm.^[2] From manufacturing and reliability perspectives, the design of TE cooler modules is often constrained by the shear stresses that result from differential thermal expansion, both during steady-state operation and during on/off cycling. In bulk systems, various strategies, such as spring-loaded systems, tension bolts, and welding, have been proposed to enhance compliance during device operation.^[3] However, none of these strategies can be directly applied to thin-film based miniaturized TE devices. Additionally, for TE devices operating at higher temperatures, a pressing need exists for stable, compliant, and low thermal resistance interface materials between the TE element and the metallic interconnects. With this motivation, we report here a scalable electrodeposition process to integrate thick-film TE materials with carbon nanotubes (CNT) arrays.

Recently, CNT arrays have been reported to exhibit excellent fatigue strength under cyclic compressive loading^[4] and as interface materials, they have been shown to achieve low thermal^[5] and electrical^[6] interface resistances at moderate contact pressures. Further, CNTs are amenable to heterogeneous integration with other materials.^[7] These attributes suggest that compliant CNT arrays can be integrated with minimal parasitic additions to the total electrical and thermal resistances of a TE device.

The common substrate for this work was 5 × 6 mm² Ni(200 nm)/Ti (800 nm)/SiO₂(1 μm)/Si(300 μm). For CNT array synthesis, some of these substrates were coated with a metal tri-layer structure of Fe (2.5 nm)/Al(10 nm)/Ti(30 nm) by electron-beam evaporation. Multi-walled CNT arrays were synthesized on these substrates by microwave-plasma chemical vapor deposition (MPCVD). The feed gases for the reaction were methane at 10 standard cubic centimeter per minute (sccm), and hydrogen at 50 sccm. The reaction temperature and pressure were 1173 K and 10 Torr (1 Torr = 133.32 Pa), respectively. Further details of the CNT synthesis by the MPCVD technique have been reported.^[5] The remaining substrates, without CNTs, served as control samples. CNT arrays were imaged using a Hitachi S-4800 field-emission scanning electron microscope (FESEM) (Fig. 1). The average length and diameter of the CNTs were 25 μm and 40 nm, respectively.

Bismuth telluride (Bi₂Te₃) was chosen as a representative TE material for TE/CNT integration by electrodeposition, as it is the parent compound of the alloys used in commercial Peltier devices optimized for cooling at temperatures near 300 K.^[8] Using a Bio-Analytical Systems (BAS) Epsilon Electrochemical System, Bi₂Te₃ was electrodeposited potentiostatically on the CNT array-coated substrates, and these films were compared with Bi₂Te₃ films electrodeposited directly on bare metallized substrates. A three-electrode setup was used with a CNT array-coated working electrode, platinum mesh counter electrode, and an Ag/AgCl (3 M NaCl, 0.175 V versus NHE) reference electrode. The deposition bath was composed of Bi³⁺ (0.75 × 10⁻² M) and HTeO₂⁺ (1 × 10⁻² M) (both Alfa Aesar, 99.999% pure) and HNO₃ (1 M) at 298 K.^[2,9] Continuous one-minute pulses at -50 mV were applied between the working and the reference electrodes for about 10 h for electrodeposition

[*] Prof. T. D. Sands, V. Rawat, P. B. Amama, K. G. Biswas
School of Materials Engineering, Purdue University West Lafayette
IN 47907 (USA)

E-mail: tsands@purdue.edu

Prof. T. D. Sands, H. Mishra, B. A. Cola, V. Rawat, P. B. Amama, K. G. Biswas, X. Xu, Prof. T. S. Fisher
Birck Nanotechnology Center 1205 West State Street, West Lafayette,
IN 47907-2057 (USA)

Prof. T. D. Sands
School of Electrical and Computer Engineering, Purdue University
West Lafayette, IN 47907 (USA)

H. Mishra, B. A. Cola, Prof. T. S. Fisher
School of Mechanical Engineering, Purdue University West Lafayette
IN 47907 (USA)

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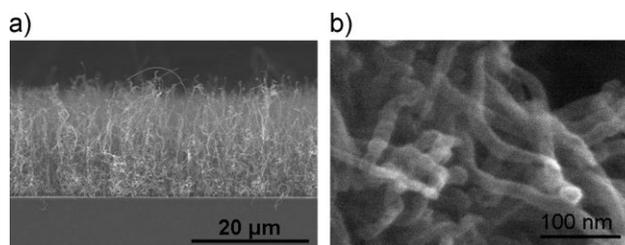


Figure 1. a) Side view of the MWCNT array and b) magnified image of MWCNT array.

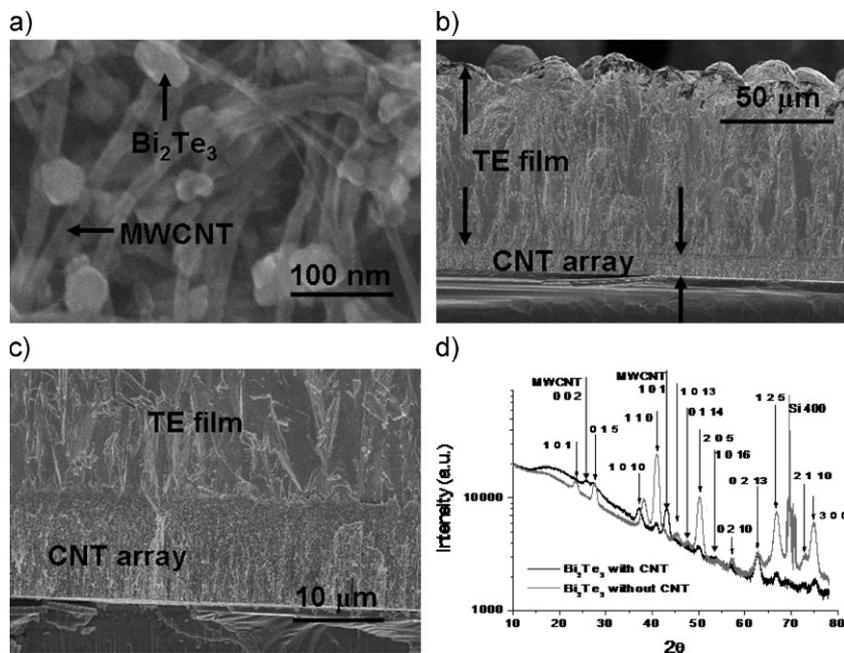


Figure 2. FESEM images showing a) nucleation of Bi₂Te₃ on individual MWCNTs, b) Bi₂Te₃ film/CNT array interface, and c) magnified view of the Bi₂Te₃ film/CNT array interface. d) Comparison of X-ray diffraction pattern obtained from Bi₂Te₃ film electrodeposited on Ni(200 nm)/Ti(800 nm)/SiO₂(1 μm)/Si(300 μm) substrate and similar Bi₂Te₃ film electrodeposited on top of CNT arrays. The background in the samples is due to the relatively small size of the samples. All of the indexed diffraction peaks correspond to Bi₂Te₃ except MWCNT 002, MWCNT 101, and Si 400 peaks, labeled accordingly.

of ~50 μm thick Bi₂Te₃ films while maintaining a current density of about 5 mA cm⁻².

FESEM images were obtained to assess the morphology of the deposited TE/CNT heterostructures in both plan and cross-sectional views (Fig. 2a–c). It can be inferred from the FESEM images that the Bi₂Te₃ electrodeposited on and around the CNTs and ultimately extended above the CNT array as a continuous polycrystalline thin film. Different durations of electrodeposition resulted in film thicknesses ranging from 1 to 100 μm. Figure 2d shows X-ray diffraction patterns obtained from a Bi₂Te₃ film deposited directly on a silicon substrate and another grown on CNT array. The X-ray diffraction patterns confirm the presence of polycrystalline Bi₂Te₃ film on the CNT array, and the crystalline quality of the film is similar to that grown under similar electrodeposition conditions directly on a metallized silicon substrate. There is a detectable influence of the CNT array on the orientation of the Bi₂Te₃ film and that effect is observed in X-ray diffraction patterns. Some of the X-ray diffraction peaks observed in the pattern from the TE/M heterostructure were found to be missing, or slightly displaced in the pattern from the TE/CNT/M heterostructure (Fig. 2d). This is expected because the CNT array underlayer, as opposed to the metal substrate, is a non-planar substrate that is mechanically compliant. Thus, any influence of the substrate on crystallographic texture or deposition-induced stress will be manifested in subtle differences in peak intensity and diffraction angle. The possibility that carbon contamination from the CNTs could have influenced the texture and lattice parameters is remote, as there was no evidence of CNT

dissolution during electrodeposition, and the Bi₂Te₃ were sufficiently thick to ensure that much of the growth occurred by Bi₂Te₃ deposition on Bi₂Te₃ rather than on CNTs directly.

For electrodeposition of Bi₂Te₃ patterns on CNT coated surfaces, towards development of a multi-coupled TE device, an optical lithography-based fabrication process was developed. On a CNT array coated surface (CNT/Ni(200 nm)/Ti(800 nm)/SiO₂(1 μm)/Si(300 μm)), MicroChemicals AZ-9260 was spin-coated to obtain a photoresist layer with a thickness of ~7 μm. The photoresist was then patterned using optical lithography to open windows for the electrodeposition of Bi₂Te₃ patterns. The underlying metallic layer beneath the CNT layer provided an electrically conducting path through the sample, and the non-conducting photoresist mold restricted electrodeposition of TE materials to the exposed CNT region. This strategy may be employed to develop p- and n-type TE legs on CNT array coated metallic surfaces for thermomechanically compliant multi-couple TE microdevices (Fig. 3a, b).

To study the effects of the CNT interface material on the relative thermomechanical robustness of the TE/CNT/M heterostructure, the temperature excursion required to induce catastrophic failure (i.e., delamination of the film) was assessed for samples with and without the CNT interface material. The samples were compared to Bi₂Te₃ films (5 μm thickness) with lateral dimensions of 5 mm by 6 mm deposited directly on either metallized (Ni(200 nm)/Ti(800 nm)/SiO₂(1 μm)/Si(300 μm)) substrates or 25 μm thick CNT arrays grown on the same metallized substrate. Samples of both types were subjected to the same thermal cycle (ramp = 600 °C h⁻¹, dwell = 1 h, quench to room temperature) in forming gas (5% H₂, 95% Ar) in a controlled environment furnace (Lindberg horizontal, three zone tube furnace).

After each thermal cycle, the films were visually analyzed for signs of cracks or peeling, and the observations are presented in Table 1. For samples without a CNT interlayer, TE films completely delaminated from the substrates for a temperature rise ($\Delta T = T_{\text{Furnace}} - T_{\text{Room}}$) greater than 200 K. However, for the

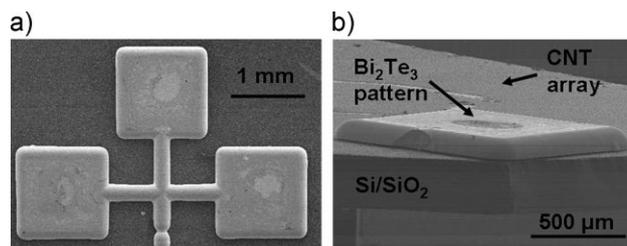


Figure 3. FESEM images showing a) top view of a Bi₂Te₃ pattern electrodeposited on top of a CNT array and b) side view of a thick Bi₂Te₃ pattern on CNT array.

Table 1. Thermomechanical failure of TE/CNT/Si and TE/Si structures.

ΔT [K]	Maximum shear displacement [μm]	TE/Si structure observations	TE/CNT/Si structure observations
50	3.12	No effect	No effect
100	6.24	No effect	No effect
150	9.36	No effect	No effect
200	12.48	Film delaminated	No effect
250	15.60	Film delaminated	No effect
300	18.72	Film delaminated	No effect
350	21.84	Film delaminated	No effect
400	24.96	Film delaminated	Cracks appeared in film

samples with CNT arrays as an interfacial layer between the Bi_2Te_3 film and the metallized substrate, the TE film remained well adhered to the substrate in all samples up to a temperature rise of 350 K (Table 1). Using the literature values for coefficients of thermal expansion for Si ($2.6 \times 10^{-6} \text{ K}^{-1}$ [10]) and Bi_2Te_3 ($13.0 \times 10^{-6} \text{ K}^{-1}$ [11]), the theoretical stress-free lateral displacement between the TE film and the substrate was calculated and is also indicated in Table 1. This maximum displacement was calculated at the perimeter of the Bi_2Te_3 film, assuming zero lateral displacement at the center. The maximum magnitude of the calculated stress-free displacement at the edge of the film with the CNT interface is $25 \mu\text{m}$ at the maximum temperature rise before failure. This value matches the average CNT length, supporting the hypothesis that the CNT interface exhibits high compliance up to the point that the CNTs are under tension. Once the CNTs are taut, further differential expansion induces mechanical failure manifested by cracking that initiates at the outer surface of the TE film under tension induced by buckling.

Effects on the thermal resistances between the metal and TE film, with and without a CNT layer, were measured as a function of interface temperature using a photoacoustic (PA) technique that has been detailed previously.^[12,13] Briefly, in the PA technique, a sinusoidally modulated fiber laser is used to periodically heat the surface of the samples. The heated area of the sample's surface is surrounded by a sealed acoustic chamber; thus, a periodic pressure signal is produced, as measured by a microphone housed in the chamber wall. The measured pressure signal is used in conjunction with a thermal model to determine thermal interface resistance.^[13]

In the PA technique, an 80 nm layer of Ti was deposited on relatively thin Bi_2Te_3 films ($\sim 30 \mu\text{m}$) to absorb the laser energy at the sample surface and to ensure adequate sensitivity to interface resistance. The measured interface resistances for the $\text{Bi}_2\text{Te}_3/\text{CNT}/\text{M}/\text{Si}$ and the $\text{Bi}_2\text{Te}_3/\text{M}/\text{Si}$ (control) samples are illustrated as a function of interface temperature in Figure 4. The thermal interface resistance for the TE/CNT structures was found to be $30 \times 10^{-6} \text{ m}^2 \text{ K W}^{-1}$, whereas for the TE/M contacts it was $\sim 2 \times 10^{-6} \text{ m}^2 \text{ K W}^{-1}$ with an uncertainty of $\pm 1 \text{ m}^2 \text{ K W}^{-1}$. From a TE device-level perspective, the interface resistance should be less than 10% of the TE leg resistance. For Bi_2Te_3 films of $100 \mu\text{m}$ thickness, the measured thermal interface resistances of CNT-TE samples is $\sim 55\%$ of the intrinsic resistance of the TE film itself (0.55 K W^{-1} , corresponding to a measured thermal conductivity of 1.8 W (m K)^{-1} at 350 K). For all control samples the interface resistance was about 3% of the TE film thermal resistance

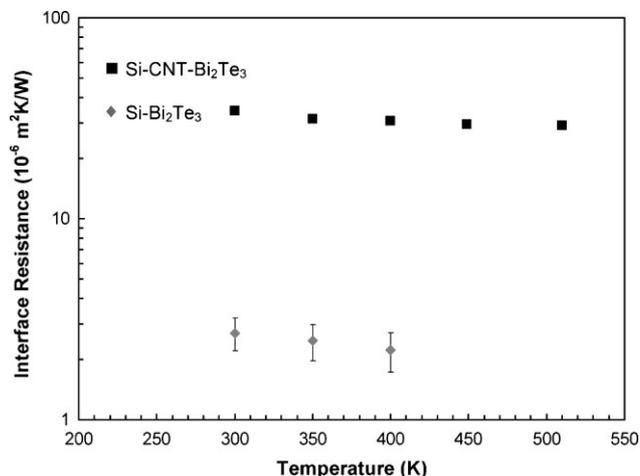


Figure 4. Thermal interface resistance as a function of interface temperature measured using the PA technique. The error in the interface resistance measurements is $\pm 1 \text{ m}^2 \text{ K W}^{-1}$ and too small to appear on the log-scale for the larger resistance values.

(0.23 K W^{-1} , corresponding to a measured thermal conductivity of 1.5 W (m K)^{-1} at 350 K). The higher thermal resistance at the TE/CNT/M/Si interface might be caused by reduction in the contact area due to the CNT array porosity. However, solid solutions of Bi_2Te_3 or other classes of TE materials with reduced thermal conductivity might still benefit from this strategy. For example, the thermal conductivity of a typical $(\text{Bi}_{0.7}\text{Sb}_{1.3})(\text{Se}_{2.91}\text{Te}_{0.09})$ alloy is 1.1 W (m K)^{-1} at 330 K.^[14] A $300 \mu\text{m}$ thick film of this TE material with the bulk value of thermal conductivity would present a thermal resistance that is about nine times larger than the interface resistance measured in the present study. Furthermore, prior work has shown that contact resistance decreases with increase in pressure on the contact.^[5,6,12] Thus, mechanical compression of the TE/CNT contacts may reduce thermal contact resistance in the CNT-TE structures.

In conclusion, integration of CNT arrays and TE materials by an electrodeposition process has been shown to improve thermomechanical compliance of the contacts. The TE/CNT/M/Si hybrid contact with a maximum lateral dimension of 6 mm showed an increase of more than 150 K in the maximum temperature excursion without mechanical failure. Thermal interface resistances of the TE/CNT/M/Si and the TE/M/Si contacts to unalloyed Bi_2Te_3 films were measured, yielding 30×10^{-6} and $2 \times 10^{-6} \text{ m}^2 \text{ K W}^{-1}$, respectively. Although the CNT interface layer increases the total thermal interface resistance, the contribution of the CNT interfacial layer to the total thermal resistance is expected to be $\sim 11\%$ of the thermal resistance of $(\text{Bi}_{0.7}\text{Sb}_{1.3})(\text{Se}_{2.91}\text{Te}_{0.09})$ alloy TE leg of $300 \mu\text{m}$ thickness. Finally, a process flow was developed for patterned synthesis of TE films on CNT array coated surfaces.

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