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# Low temperature CVD of thermoelectric SnTe thin films from the single source precursor, [<sup>n</sup>Bu<sub>3</sub>Sn(Te<sup>n</sup>Bu)]<sup>†</sup>

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This work has demonstrated that the single source precursor [<sup>n</sup>Bu<sub>3</sub>Sn(Te<sup>n</sup>Bu)], bearing *n*-butyl groups and containing the necessary 1 : 1 Sn : Te ratio, facilitates growth of continuous, stoichiometric SnTe thin films. This single source CVD precursor allows film growth at significantly lower temperatures (355–434 °C at 0.01–0.05 Torr) than required for CVD from SnTe powder. This could be advantageous for controlling the surface states in topological insulators. The temperature-dependent thermoelectric performance of these films has been determined, revealing them to be p-type semiconductors with peak Seebeck coefficient and power factor values of 78 μV K<sup>-1</sup> and 8.3 μW K<sup>-2</sup> cm<sup>-1</sup>, respectively, at 615 K; comparing favourably with data from bulk SnTe. Further, we have demonstrated that the precursor facilitates area selective growth of SnTe onto the TiN regions of SiO<sub>2</sub>/TiN patterned substrates, which is expected to be beneficial for the fabrication of micro-thermoelectric generators.

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## Introduction

With escalating concerns regarding climate change and a rapidly increasing demand for wireless power generation for sensors and other applications, thermoelectric materials are emerging as a key to advancement in both environmental and technological fields.<sup>1,2</sup> Thermoelectric materials enable the conversion between heat and electricity and thus allow for both active cooling systems and the recycling of otherwise wasted heat energy.<sup>3,4</sup> These processes rely on the Peltier and Seebeck effects, which are the conversion from electricity to cooling and the conversion of a heat gradient to the flow of charge, respectively.<sup>5</sup> The materials that perform best for thermoelectric applications have a combination of high electrical conductivity, a large Seebeck coefficient and a low thermal conductivity. A useful, but not to be used alone, comparative value is the unitless figure of merit, *ZT*, which is described by the equation  $ZT = \frac{S^2 \sigma T}{\kappa}$ , where the Seebeck coefficient, *S*, the

electrical conductivity,  $\sigma$ , the average temperature across the material, *T* and the thermal conductivity of the material,  $\kappa$ , are all brought together.<sup>6</sup>

One particular group of semiconducting materials, the metal chalcogenides, including the tellurides, shows great potential for thermoelectric applications. In particular, Bi<sub>2</sub>Te<sub>3</sub> and Sb<sub>2</sub>Te<sub>3</sub> are among the most promising thermoelectric materials for near room temperature applications, while PbTe has been used for mid-temperature applications, including for power generation in space, since the 1960s.<sup>5,7</sup> However, due to the high toxicity of lead, alternative materials are desirable. A potential replacement for PbTe is SnTe, as it possesses many similarities to PbTe in both its crystal and electronic structures.<sup>8,9</sup> SnTe exhibits an intrinsically high charge carrier concentration, which results in a relatively low Seebeck coefficient,<sup>10</sup> but optimisation of the material through doping and alloying mean that SnTe offers great promise for thermoelectric applications.

SnTe has also gained significant interest due to the exciting properties relevant to other applications that it displays. Key examples are as a topological crystalline insulator,<sup>11,12</sup> IR detection and radiation receivers<sup>13,14</sup> and in near IR photovoltaics.<sup>15</sup>

Most previous work on the thermoelectric properties of SnTe relates to the bulk form, but given the low natural abundance of tellurium, the sustainability of bulk tellurium-based thermoelectrics is questionable. On the other hand, the quantity of material required for thin film based thermoelectric microgenerators is very considerably reduced (by >95%), leading to the prospect of using this in internet of things (IoT) based applications. While physical vapour deposition (PVD)<sup>14</sup>

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mass. Using these the final equation gives the vapour pressure,  $p$ . At each of the temperatures measured the value of  $k$  was first calculated using the experimentally determined value for  $\nu$  and the known value for  $p$ .<sup>29</sup> Using the value for  $k$  gained in this method, the vapour pressure of the single source precursor could be calculated.

### Synthesis of [<sup>n</sup>Bu<sub>3</sub>Sn(Te<sup>n</sup>Bu)]

Tellurium granules (2.846 g,  $2.23 \times 10^{-2}$  mol) were ground into a fine powder under an inert atmosphere in the glovebox before being suspended in THF (30 mL). The suspension was then frozen in liquid nitrogen (77 K) before the dropwise addition of <sup>n</sup>BuLi solution in hexanes (13 mL,  $2.18 \times 10^{-2}$  mol). After the mixture had slowly thawed, the solution initially turned yellow and then red. Further <sup>n</sup>BuLi solution (0.3 mL,  $5.03 \times 10^{-4}$  mol) was then added dropwise until a pale yellow colour persisted. A solution of <sup>n</sup>Bu<sub>3</sub>SnCl (6.856 g,  $2.11 \times 10^{-3}$  mol) in THF (30 mL) was then added dropwise, while it was cooled using an ice bath. The reaction mixture was stirred vigorously for 24 h, leaving an orange solution. The volatiles were then removed *in vacuo* and the orange oily residue was then washed twice with hexane (30 mL), leading to the precipitation of LiCl. This was removed by cannula filtration before the filtrate was taken to dryness *in vacuo*, leaving an orange oil. The compound is somewhat moisture sensitive and was therefore handled and stored under an inert atmosphere to avoid degradation, although it can be handled briefly in air without degradation. Yield 8.608 g, 86%. Elemental analysis: calc. for C<sub>16</sub>H<sub>36</sub>SnTe (474.77): C 40.48, H 7.64%. Found: C 40.60, H 8.65%. <sup>1</sup>H NMR (CDCl<sub>3</sub>):  $\delta$ /ppm = 0.90 (two overlapping t,  $J$  = 8.00 Hz, [12H]), 1.19 (t with <sup>119</sup>Sn satellites, [6H]), 1.34 (overlapping m, [8H]), 1.55 and 1.65 (overlapping m, [8H]), 2.52 (t with overlapping <sup>125</sup>Te satellites, [2H]). <sup>13</sup>C{<sup>1</sup>H} NMR (CDCl<sub>3</sub>):  $\delta$ /ppm = -7.05 (<sup>1</sup>J<sub>125Te13C</sub> = 133 Hz, <sup>2</sup>J<sub>119Sn13C</sub> = 13 Hz, CH<sub>2</sub>), 12.54 (<sup>1</sup>J<sub>119Sn13C</sub> = 299 Hz, <sup>1</sup>J<sub>117Sn13C</sub> = 286 Hz, CH<sub>2</sub>), 13.27 (CH<sub>3</sub>), 13.61 (CH<sub>3</sub>), 25.01 (CH<sub>2</sub>), 26.96 (<sup>2</sup>J<sub>119Sn13C</sub> = 63 Hz, CH<sub>2</sub>), 29.69 (<sup>3</sup>J<sub>119Sn13C</sub> = 21 Hz, CH<sub>2</sub>), 37.34 (<sup>3</sup>J<sub>125Te13C</sub> = 8 Hz CH<sub>2</sub>). <sup>119</sup>Sn{<sup>1</sup>H} NMR (CD<sub>2</sub>Cl<sub>2</sub>/CH<sub>2</sub>Cl<sub>2</sub>):  $\delta$ /ppm = -15.62 (<sup>1</sup>J<sub>119Sn125Te</sub> = 2765 Hz). <sup>125</sup>Te{<sup>1</sup>H} NMR (CD<sub>2</sub>Cl<sub>2</sub>/CH<sub>2</sub>Cl<sub>2</sub>):  $\delta$ /ppm = -536.51 (<sup>1</sup>J<sub>119Sn125Te</sub> = 2704 Hz).

### Thin film growth by low pressure CVD onto fused silica substrates

The fused silica substrates (8 mm × 20 mm) were washed with deionised water, followed by ethanol, before being dried thoroughly in an oven before use. In a LPCVD experiment the precursor is loaded (5–40 mg) into a closed end silica tube (1 cm diameter, 45 cm long). With the tube held horizontally,



Scheme 1 Synthesis route to [<sup>n</sup>Bu<sub>3</sub>Sn(Te<sup>n</sup>Bu)].

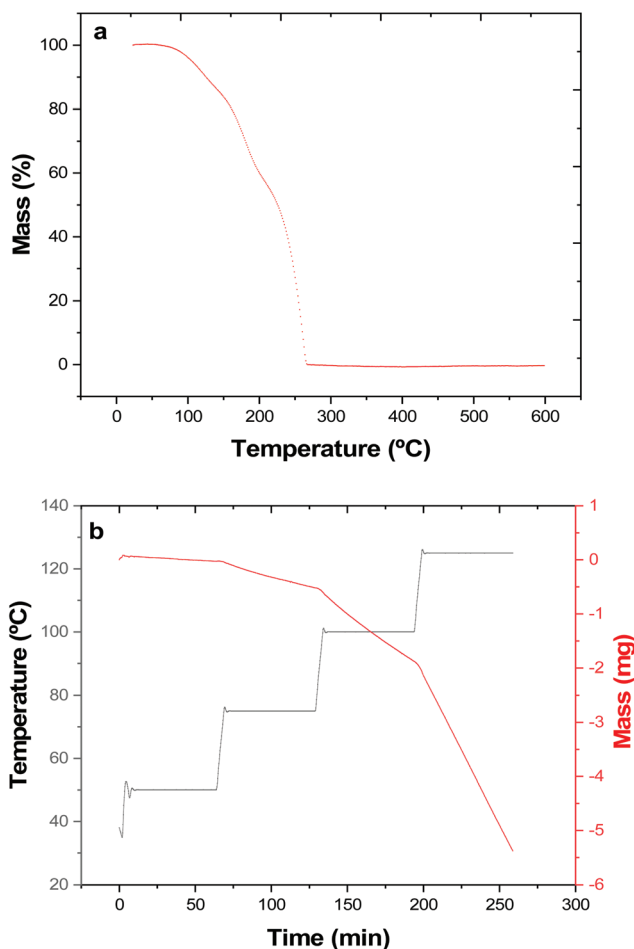


Fig. 1 (a) Thermogravimetric analysis of [<sup>n</sup>Bu<sub>3</sub>Sn(Te<sup>n</sup>Bu)] showing the percentage mass loss with temperature at a ramp rate of 5 °C min<sup>-1</sup> and a constant flow of argon gas. (b) Isothermal step TGA experiment of [<sup>n</sup>Bu<sub>3</sub>Sn(Te<sup>n</sup>Bu)] used for vapour pressure determination.

Table 1 Vapour pressures of [<sup>n</sup>Bu<sub>3</sub>Sn(Te<sup>n</sup>Bu)] determined as a function of temperature. Boiling point,  $T_{\text{vap}}$ , from Sigma Aldrich SDS and estimated from TGA data,<sup>a</sup> enthalpy of vaporisation calculated using the Clausius–Clapeyron equation<sup>b</sup> and reference values given for <sup>n</sup>Bu<sub>4</sub>Sn, as well as the entropy of vaporisation calculated using  $\Delta H_{\text{vap}}/T_{\text{vap}}$  along with reference values for <sup>n</sup>Bu<sub>4</sub>Sn

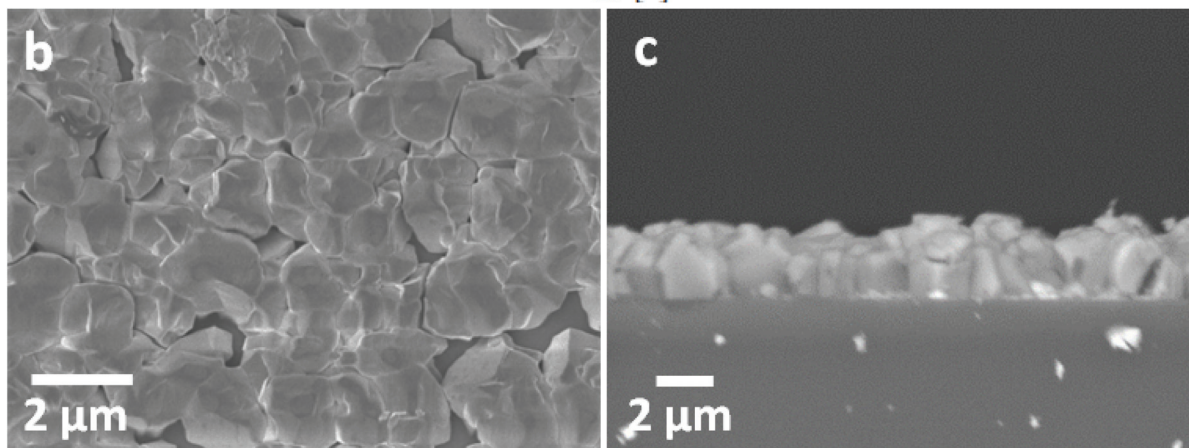
Temperature (K)	<sup>n</sup> Bu <sub>4</sub> Sn vapour pressure (Pa) <sup>36</sup>	[ <sup>n</sup> Bu <sub>3</sub> Sn(Te <sup>n</sup> Bu)] vapour pressure (Pa)
323.15	17	3 ± 1
348.15	70	23 ± 4
373.15	240	80 ± 13
398.15	703	233 ± 37
$T_{\text{vap}}$	518.15 K	473.15 K <sup>a</sup>
$\Delta H_{\text{vap}}$	53 kJ mol <sup>-1</sup> (67.8 kJ mol <sup>-1</sup> )	67 kJ mol <sup>-1</sup> ± 15
$\Delta S_{\text{vap}}$	100 J K <sup>-1</sup> mol <sup>-1</sup>	140 J K <sup>-1</sup> mol <sup>-1</sup> ± 32
	(130 J K <sup>-1</sup> mol <sup>-1</sup> )	

this was followed by the substrates placed next to one another (lengthways along the tube starting closest to the precursor bulb). The assembly of the CVD tube is performed under an inert atmosphere inside a glovebox before being moved into



position in a tube furnace, insulated using silica wool at both ends, and being placed under a constant reduced pressure (0.01–0.05 mmHg). The Lenton (LTF 12/50/300) tube furnace was then brought to temperature (furnace set temperature = 475 °C; substrate temperature where deposition observed = 355 to 434 °C). Once the desired temperature had been reached the tube was repositioned to bring the precursor bulb closer to the hot zone so that volatilisation could commence (the temperature at the precursor bulb was between 274 to 355 °C, with

a rate of mass transfer of between 1.72 and 3.09 mg min<sup>-1</sup>). The time taken for the deposition to occur once the tube furnace was at temperature was typically 5–18 min (the rates of growth were between 64 and 125 nm min<sup>-1</sup>). After cooling and letting down to a dinitrogen atmosphere, the CVD tube was transferred back into the glovebox where the substrates onto which the films had deposited were removed and stored. The low pressure CVD experiments produced light grey films of SnTe with typical coverage of 1–4 cm × 0.8 cm.



**Fig. 2** (a) Grazing incidence XRD pattern for SnTe thin film (thickness  $\sim 2 \mu\text{m}$ ) with XRD pattern for bulk cubic SnTe ( $Fm\bar{3}m$ ) (red).<sup>8</sup> The broad feature observed around  $21^\circ$  is due to the fused silica substrate beneath the SnTe film. Top down SEM (b) and cross-sectional SEM (c) of a SnTe film.





### Thin film growth by low pressure CVD onto patterned SiO<sub>2</sub>/TiN substrates

TiN/SiO<sub>2</sub> patterned substrates (8 mm × 20 mm) were prepared as described previously.<sup>26</sup> These were loaded lengthways into a closed end silica tube containing <3 mg of the [<sup>n</sup>Bu<sub>3</sub>Sn(Te<sup>n</sup>Bu)] precursor. The deposition process parallels that onto the unpatterned substrates, with the subtle differences of the lower precursor loading and the position of the tube causing the slower volatilisation of the precursor (and hence better discrimination of the TiN vs. SiO<sub>2</sub> surfaces), thus the duration of the depositions is longer (30–45 min).

### Thin film characterisation

Grazing incidence X-ray diffraction (XRD) patterns were collected using a Rigaku SmartLab system (Cu-Kα, λ = 1.5418 Å) with parallel X-ray beam and a HyPix detector in 1D mode. A 2θ scan range of 10–80° and an incident angle (ω-offset) of 1° were used. The crystalline phase of the film was determined by comparison with a literature XRD pattern, accessed *via* the Inorganic Crystal Structure Database (ICSD),<sup>31</sup> and lattice parameters were obtained by further optimisation of the fit using PDXL.<sup>32</sup>

The scanning electron microscope used was a FEI XL30 ESEM with a tungsten filament electron source and ThermoFisher UltraDry EDX, 10 mm<sup>2</sup> detector. The SEM images were all captured at magnifications of 2000, 5000 and 10 000 times at accelerating voltages of either 10 or 15 keV. The Raman spectra were obtained using a Renishaw inVia confocal Raman microscope with a 785 nm laser. The incident laser power was adjusted to 2.5 mW for all samples.

### Thermoelectric and electrical property measurements

Variable temperature resistivity, carrier concentration, and carrier mobility were determined by Hall measurements (Nanometrics HL5500PC) from 125 to 450 K in steps of 25 K under a magnetic field of 0.5 T. Care was taken to ensure linear contact between the probes and sample before each measurement and the current was optimised to ensure normal Ohmic conduction with maximum voltage signal. Variable temperature Seebeck coefficient and electrical resistivity were measured using the Joule Yacht MRS-3L from 300 to 600 K in steps of 25 K. The Seebeck coefficient was measured using the differential method with a maximum temperature difference of 10 K. The resistivity was measured using a 4-point probe method and corrected for geometric error using the results from the Hall measurement system.

## Results and discussion

### Synthesis and characterisation of the [<sup>n</sup>Bu<sub>3</sub>Sn(Te<sup>n</sup>Bu)] precursor

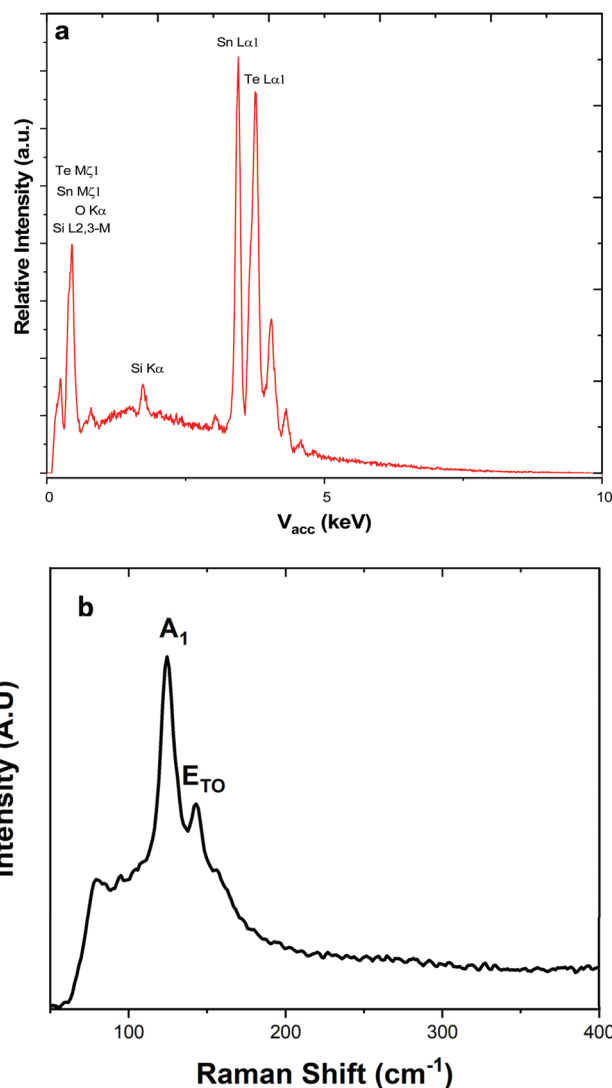
The organometallic tin(IV) compound, [<sup>n</sup>Bu<sub>3</sub>Sn(Te<sup>n</sup>Bu)], was selected as a potentially viable single source precursor for the deposition of SnTe thin films since the *n*-butyl substituents on both the tin and tellurium provide a clean and low energy

decomposition pathway *via* β-hydride elimination, the 1 : 1 Sn : Te ratio in the precursor should aid deposition of the targeted binary phase and avoid incorporation of excess Te, while the volatility of the precursor (oil) was anticipated to be well-suited to a low pressure CVD procedure.<sup>33</sup> Previous work has

**Table 2** EDX data for four SnTe thin film samples (atomic percentages)

Tile	O% <sup>a</sup>	Si%	Sn%	Te%
Deposition 1 Tile 1	16.4	2.1	41.4	40.1
Deposition 2 Tile 1	16.9	2.3	41.1	39.8
Deposition 2 Tile 2	15.8	10.7	41.9	41.6
Deposition 3 Tile 1	23.5	2.3	37.5	37.7

<sup>a</sup> Note that the %O arises from the substrate and is exaggerated due to overlap with peaks characteristic of Sn and Te.



**Fig. 3** (a) EDX spectrum highlighting the elements present within the SnTe film, the silicon present is attributed to the substrate. (b) Raman spectrum with the data highlighting the A<sub>1</sub> and E modes at 124 and 143 cm<sup>-1</sup>, respectively.<sup>1</sup>





line films comprised of block-like particles. The cross-sectional SEM analysis determined the film thickness to be around  $\sim 1\text{--}2\ \mu\text{m}$  and the grain size is around  $1\ \mu\text{m}$ .<sup>3</sup>

Energy dispersive X-ray analysis showed that all the films had compositions within experimental error of stoichiometric SnTe (Table 2). Peaks corresponding to Si and O from the underlying substrates were present in a few samples (indicating thinner films), but no impurities were evident. This was also supported by Raman scattering, which showed only

features expected for SnTe, with the  $A_1$  and E modes at  $124$  and  $143\ \text{cm}^{-1}$ , respectively (Fig. 3), and with no contamination evident.<sup>11,37</sup>

### Thermoelectric measurements

In order to evaluate the thermoelectric performance of the SnTe thin films deposited *via* our CVD process, the temperature-dependent Hall and Seebeck measurements were conducted. The as-deposited SnTe film exhibits a metallic trans-

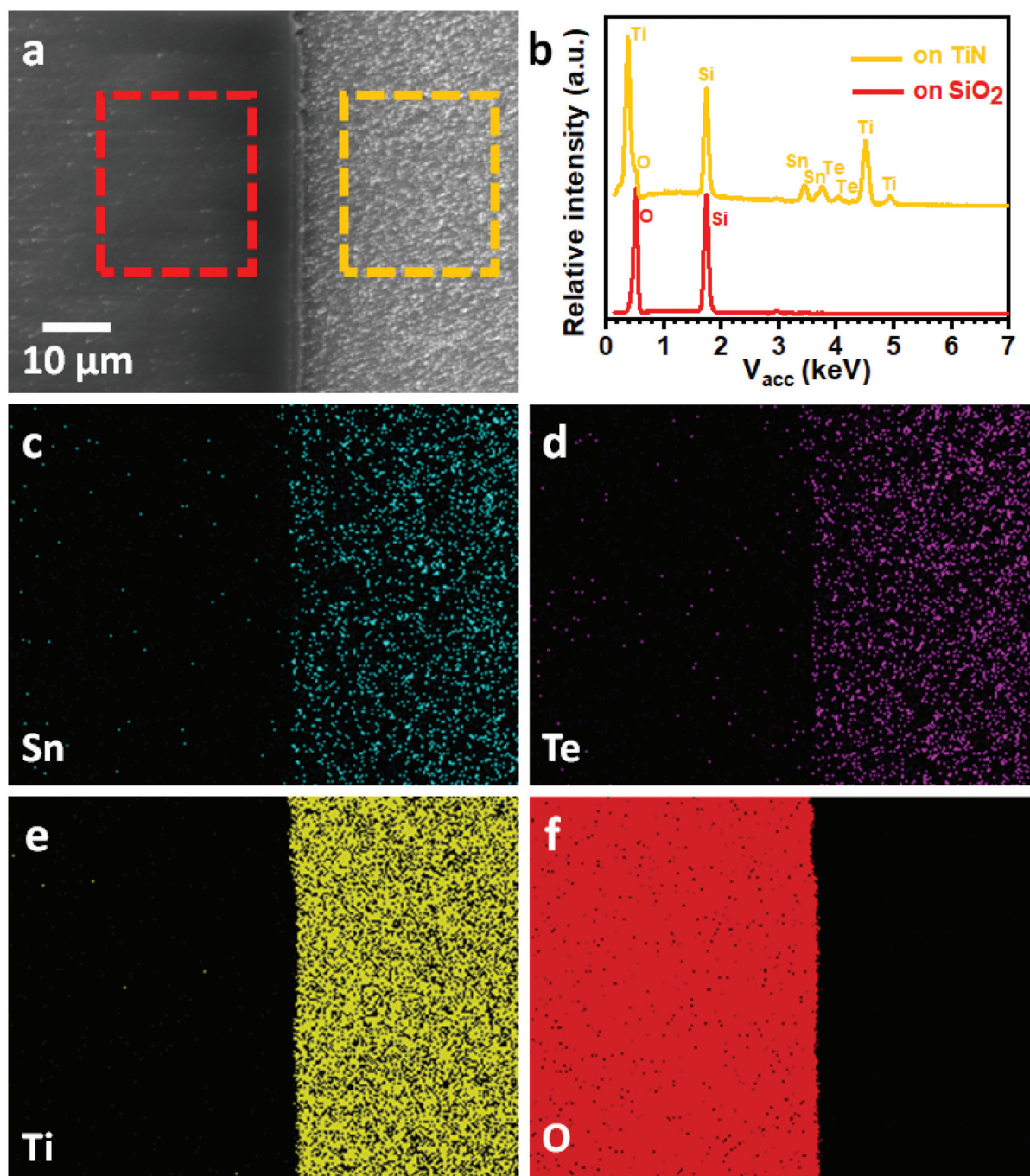


Fig. 5 (a) Top-view SEM image of a TiN/SiO<sub>2</sub> patterned substrate with SnTe selectively deposited on the TiN surface (right). (b) EDX spectra taken from the rectangular areas indicated in (a). EDX element mapping of (c) Sn, (d) Te, (e) Ti and (f) O.





portation behaviour with room temperature electrical conductivity of  $2434 \text{ S cm}^{-1}$  (Fig. 4a). This high electrical conductivity can be ascribed to the intrinsic Sn vacancy induced p-type carrier concentration.<sup>10</sup> This is confirmed by the Hall measurement results where high p-type carrier concentrations in the range between  $(4\text{--}5) \times 10^{21} \text{ cm}^{-3}$  were observed, as shown in Fig. 4b. While this carrier concentration remains relatively constant, the mobility decreases from  $ca. 5 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$  to  $3 \text{ cm}^2 \text{ V}^{-1} \text{ s}^{-1}$  as the temperature increases from 100 K to 450 K. This is likely to be caused by the stronger lattice scattering at elevated temperatures, resulting in the reduction of electrical conductivity.

Fig. 4c presents the temperature dependent Seebeck coefficient of the as-deposited SnTe thin film. The p-type conductivity is further confirmed by the positive Seebeck coefficient values obtained. The Seebeck coefficient for the as-deposited SnTe film is  $ca. 12 \mu\text{V K}^{-1}$  at room temperature and is increased to  $ca. 78 \mu\text{V K}^{-1}$  at higher temperature (615 K). The power factor of the SnTe film is shown in Fig. 4d, with a peak value of  $8.3 \mu\text{W K}^{-2} \text{ cm}^{-1}$  at 615 K. These results are comparable to the other reported values for undoped SnTe samples,<sup>10,38–40</sup> also confirming the high quality of the SnTe films produced from the single source precursor used in this work. It is worth mentioning that high hole carrier concentration and large energy separation ( $ca. 0.3\text{--}0.4 \text{ eV}$ ) between light hole and heavy hole valence bands have limited the thermoelectric properties of un-doped SnTe.<sup>41</sup> Significant improvements in the Seebeck coefficient of bulk SnTe have been achieved by pioneering works using doping/alloying to induce valence band convergence.<sup>40–42</sup> We have also demonstrated recently the capability of alloying  $\text{Bi}_2\text{Te}_3$  thin films with both  $\text{Bi}_2\text{Se}_3$  and  $\text{Sb}_2\text{Te}_3$  using single source precursors.<sup>43</sup> Future work will explore using a similar strategy to further improve the thin film SnTe thermoelectric performance.

### Selective deposition of SnTe onto patterned substrates

We have previously demonstrated selective deposition for a range of metal chalcogenide materials onto specific regions of lithographically patterned substrates using tailored single source precursors.<sup>26,27</sup> A correlation between the electrical conductivity of the film and the selective deposition performance has been established, with higher conductivity leading to enhanced selectivity.<sup>41</sup> We have also shown that deposition of the highly conductive SnTe onto TiN/SiO<sub>2</sub> patterned substrates in this work results in selective growth on the TiN surface in preference to the SiO<sub>2</sub> surface, as shown Fig. 5a. EDX spectra (Fig. 5b) on the TiN surface (yellow rectangular area in Fig. 5a) clearly shows the existence of Sn and Te, while only Si and O peaks can be observed from the EDX data taken on the SiO<sub>2</sub> surface (red rectangular area in Fig. 5a). This selective deposition behaviour is further demonstrated by EDX element mapping (Fig. 5c–f) where the Sn and Te are almost entirely confined to the TiN areas of the pattern. It is worth mentioning that the selective deposition behaviour allows great positional control over the material growth and is beneficial for the fabrication of micro-thermoelectric generators.<sup>44</sup>

## Conclusions and outlook

This work has demonstrated that the single source precursor [ $^n\text{Bu}_3\text{Sn}(\text{Te}^n\text{Bu})$ ], bearing *n*-butyl groups and containing the required 1 : 1 Sn : Te ratio, facilitates low pressure CVD growth of continuous, stoichiometric SnTe thin films very effectively. The temperature-dependent thermoelectric performance of these films confirms them to be p-type semiconductors with peak Seebeck coefficient and power factor values of  $78 \mu\text{V K}^{-1}$  and  $8.3 \mu\text{W K}^{-2} \text{ cm}^{-1}$ , respectively, at 615 K, comparing favourably with data from bulk SnTe. We have also demonstrated that this precursor facilitates selective growth of SnTe onto the TiN regions of SiO<sub>2</sub>/TiN patterned substrates, which is expected to be beneficial for the fabrication of micro-thermoelectric generators.

This single source precursor also allows for SnTe film growth at a significantly lower temperature than that required for CVD from SnTe powder, which is expected to be advantageous for controlling the surface states in topological insulators.

## Conflicts of interest

The authors have no conflicts to declare.

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