

# Materials Advances

rsc.li/materials-advances



ISSN 2633-5409

**PAPER**

Chitra Gurnani *et al.*  
Solution-based *in situ* deposition of  $\text{Sb}_2\text{S}_3$  from a single  
source precursor for resistive random-access memory  
devices

Cite this: *Mater. Adv.*, 2023,  
4, 4119Received 1st May 2023,  
Accepted 1st August 2023

DOI: 10.1039/d3ma00205e

rsc.li/materials-advances

# Solution-based *in situ* deposition of Sb<sub>2</sub>S<sub>3</sub> from a single source precursor for resistive random-access memory devices†

Sayali Shrishail Harke,<sup>a</sup> Tongjun Zhang,<sup>b</sup> Ruomeng Huang<sup>b</sup> and Chitra Gurnani<sup>ib\*</sup>

A one-step, simple, scalable, reproducible, low-temperature, and *in situ* solvothermal deposition method has been established for the growth of Sb<sub>2</sub>S<sub>3</sub> on FTO using [Sb(S<sub>2</sub>P(O)(Pr)<sub>2</sub>)<sub>3</sub>] as a single source precursor without a binding agent. XRD, Raman, SAED, and HRTEM results revealed the crystalline orthorhombic stibnite phase. The sheaf-like Sb<sub>2</sub>S<sub>3</sub> exhibited a band gap energy of 1.72 eV. The Sb<sub>2</sub>S<sub>3</sub> film is uniform and well-adhered and is further developed as a novel resistive random-access memory material. The Ag/Sb<sub>2</sub>S<sub>3</sub>/FTO memristive device demonstrated low operating voltage and the performance of the device over multiple cycles revealed dependable bipolar resistive switching behaviour and an ON/OFF ratio of ca. 10.

## 1. Introduction

A promising aspect of new non-volatile memory technologies is their predicted capacity gains, non-volatility, and lack of idle energy. The most promising alternative is resistive random access memory (RRAM), which is based on resistive switching (RS). RRAM devices have recently undergone rapid development, due to their unique qualities, including consumption of low power, fast switching rates, superior scalability, and increased data endurance and retention capabilities.<sup>1–10</sup> Developing highly effective and low-powered devices is important for RRAM since it must satisfy high technological needs with extensive integration. However, it is difficult to achieve an effective performance of the device within a single device with low switching voltage, operating current, a better ON/OFF ratio, and dependable switching behaviour.<sup>11</sup> Resistive switching has been detected in oxides,<sup>12</sup> chalcogenides,<sup>13</sup> and organic,<sup>14</sup> as well as organic–inorganic hybrid materials.<sup>15,16</sup>

Metal oxides (such as WO<sub>3</sub>, SiO<sub>2</sub>, ZrO<sub>2</sub>, TiO<sub>2</sub>, and CuO) and germanium chalcogenides (Ge<sub>x</sub>S<sub>x</sub> or Ge<sub>x</sub>Se<sub>x</sub>, and Ge<sub>x</sub>Te<sub>y</sub>) exhibit attractive switching characteristics.<sup>17–23</sup> There is a constant

need to explore novel memristive materials. Sb<sub>2</sub>S<sub>3</sub> is an efficient light-absorbing material and has been reported to exhibit light induced transformation of resistive switching polarity in Sb<sub>2</sub>S<sub>3</sub> based organic–inorganic hybrid devices. Moreover, Sb<sub>2</sub>S<sub>3</sub> possesses a high absorption coefficient, suitable bandgap (1.7 eV), earth-abundance, high stability, reduced toxicity,<sup>24–30</sup> high resistance and low carrier concentration.<sup>31,32</sup> To our knowledge, there have been only two reports, where Sb<sub>2</sub>S<sub>3</sub> was used for memristor applications.<sup>33,34</sup> Consequently, there is a lot of interest in the development of efficient and novel routes to synthesize Sb<sub>2</sub>S<sub>3</sub>.

The majority of the published investigations on the deposition of Sb<sub>2</sub>S<sub>3</sub> emphasized on the utilization of dual-source precursors (DSPs), and very few studies exploited the usage of single-source precursor (SSP) complexes. SSPs can increase atom efficiency and simplicity of handling while enabling precursor design to control characteristics like stoichiometry and morphology.<sup>35</sup> The SSP methods offer mildness, safety, and a more streamlined production process compared to the usage of DSP methods, which necessitates strict control over stoichiometry.<sup>36</sup> Only high temperature routes such as aerosol-assisted chemical vapor deposition (AACVD),<sup>37,38</sup> and the electrodeposition method<sup>34</sup> have been explored for Sb<sub>2</sub>S<sub>3</sub>; however, low temperature and solution processible routes using SSPs for deposition of Sb<sub>2</sub>S<sub>3</sub> have not been explored so far. Solvothermal deposition is simple in operation, a low-temperature deposition process, controls the overall growth rate, doesn't involve any complicated devices, has low production cost, and has the prospect of being suitable for large-scale industrial production. Sb<sub>2</sub>S<sub>3</sub> (stibnite) powder has been generated *via* a solvothermal

<sup>a</sup> Ecole Centrale School of Engineering, Mahindra University (MU), Hyderabad, India. E-mail: chitra.gurnani@mahindrauniversity.edu.in

<sup>b</sup> School of Electronics and Computer Science, University of Southampton, Southampton, UK

† Electronic supplementary information (ESI) available: Original IR, <sup>1</sup>H, and <sup>31</sup>P{<sup>1</sup>H} NMR spectroscopic data for [Et<sub>3</sub>NH][[(OC<sub>3</sub>H<sub>7</sub>)<sub>2</sub>S<sub>2</sub>P] and [Sb(S<sub>2</sub>P(O)(Pr)<sub>2</sub>)<sub>3</sub>], the size distribution histogram and cross-sectional view of deposited Sb<sub>2</sub>S<sub>3</sub> on the FTO substrate and bipolar non-volatile resistive switching behaviours plot of the W/Sb<sub>2</sub>S<sub>3</sub>/FTO. See DOI: <https://doi.org/10.1039/d3ma00205e>



method<sup>39,40</sup> with high boiling solvent oleylamine and *via* a refluxing route.<sup>41</sup> Nonetheless, *in situ* solvothermal deposition of Sb<sub>2</sub>S<sub>3</sub> thin films from SSPs without any binding agents and additional step of annealing has lingered unexplored. Dithiophosphate (DTP) complexes exhibit diverse coordination patterns.<sup>42</sup> They are well defined molecular species and contain performed M–S bonds that decompose when heated, either in a vapor or solution phase, with the easy loss of organics and creation of low-dimensional metal sulfides as a result. The incredible adaptability of DTP complexes as SSPs is due to their simplicity in synthesis and handling, long metal complex shelf lives, and relative stability. To date, there has been no report on the deposition of Sb<sub>2</sub>S<sub>3</sub> films using SSPs from DTP by an *in situ* solvothermal process. Consequently, there is immense potential to explore novel routes for mass manufacturing of Sb<sub>2</sub>S<sub>3</sub> films towards memristors.

In this study, we report a one-step, simple, scalable, reproducible, and *in situ* solvothermal deposition for the growth of Sb<sub>2</sub>S<sub>3</sub> on a substrate. The deposition of Sb<sub>2</sub>S<sub>3</sub> was performed at a very low deposition temperature by using SSP [Sb{S<sub>2</sub>P{O(Pr)}<sub>2</sub>}<sub>3</sub>] without a binding agent. The deposited film is uniform and well-adhered, and further utilized as a resistive random-access memory device, which demonstrated typical bipolar resistive switching behaviour. Remarkably, the Ag/Sb<sub>2</sub>S<sub>3</sub>/FTO memristive device exhibited characteristics of low operating voltage, stability over multiple cycles and an ON/OFF ratio of *ca.* 10.

## 2. Experimental

### 2.1 Materials

SbCl<sub>3</sub> and P<sub>2</sub>S<sub>5</sub> were procured from Sigma Aldrich and utilized as received. Propan-1-ol (Pr-99%) (b.p. 97 °C), and toluene (99.5%) (b.p. 110.6 °C) were bought from SDFCL and FINAR, respectively. Triethylamine (99%), *n*-hexane (99%), dichloromethane (DCM) (99.5%), ethanol (99.9%), and ethylene glycol (99.5%) were purchased from SDFCL, SRL, FINAR, Analytical reagent, and Fischer Scientific, respectively, and used as received.

### 2.2 Preparation of precursor and characterization

All the reactions were performed in Schlenk and vacuum lines. *n*-Hexane, toluene, propan-1-ol, dichloromethane and triethylamine were freshly distilled prior to use. Ethanol and ethylene glycol were used as received. On a Nicolet iS5 Thermo Scientific equipment, IR spectra were captured using KBr discs in the wavelength range of 4000–400 cm<sup>-1</sup>. On a Bruker Avance Neo 500 MHz NMR spectrometer, <sup>1</sup>H} NMR and <sup>31</sup>P{<sup>1</sup>H} NMR spectra were collected in CDCl<sub>3</sub> solution at room temperature in SAIF, Punjab University, Chandigarh.

[Et<sub>3</sub>NH][{(OC<sub>3</sub>H<sub>7</sub>)<sub>2</sub>S<sub>2</sub>P}]. P<sub>2</sub>S<sub>5</sub> (2.5 g, 1.12 mmol) was gradually dropped and stirred with distilled toluene (20 mL). Next, freshly distilled propanol (2.7 g, 4.48 mmol) was poured to the P<sub>2</sub>S<sub>5</sub> suspension in toluene. The solution was stirred until it dissolves completely at 80–90 °C. The solution was then allowed to cool to room temperature and Et<sub>3</sub>N (3.2 mL, 2.24 mmol) was dropped into the aforementioned solution. The precipitate was

not observed after the addition of Et<sub>3</sub>N, and *n*-hexane was added to the above solution to assist precipitation. A white coloured precipitate was obtained which was filtered and dried in a vacuum. Furthermore, the precipitate was dissolved in DCM and layered with *n*-hexane for recrystallization, resulting in colourless crystals standing after 3–4 days.

[Et<sub>3</sub>NH][{(OC<sub>3</sub>H<sub>7</sub>)<sub>2</sub>S<sub>2</sub>P}]. Yield: (6.3 g, 88.2%); anal: calcd M.W. of C<sub>12</sub>H<sub>30</sub>O<sub>2</sub>PS<sub>2</sub>N: 315.2; <sup>1</sup>H} NMR (CDCl<sub>3</sub>, 298 K, δ ppm): 0.95 (t, [6H], –CH<sub>3</sub> of propanol), 1.42 (t, [9H], –CH<sub>3</sub> of Et<sub>3</sub>N), 1.7 (m, [4H], –CH<sub>2</sub>CH<sub>3</sub> of propanol), 3.3 (m, [4H], –CH<sub>2</sub> of propanol), 3.95 (m, [6H], –CH<sub>2</sub> of Et<sub>3</sub>N); <sup>31</sup>P{<sup>1</sup>H} (CDCl<sub>3</sub>, 298 K, δ ppm): 110.66(s); IR (KBr) (cm<sup>-1</sup>): 2967(s), ν(N–H); 1120(s), ν(P–O–C); 832(s), ν(P–O–C); 675(s), ν(P=S) and 555(s), ν(P–S).

[Sb{S<sub>2</sub>P{O(Pr)}<sub>2</sub>}<sub>3</sub>]. A suspension of {[Et<sub>3</sub>NH][{(OC<sub>3</sub>H<sub>7</sub>)<sub>2</sub>S<sub>2</sub>P}]}<sub>2</sub> (0.85 g, 2.7 mmol) in toluene (10 ml) was typically added to antimony trichloride (0.21 g, 0.9 mmol) in toluene (10 ml) solution with a 3:1 molar ratio and continuously stirred at room temperature for 1 h. Filtration under anhydrous conditions was performed to extract the triethylamine chloride precipitate, and excess solvent has been eliminated from the filtrate *in vacuo* to produce a light pale yellow precipitate. To produce yellow needle-shaped crystals, the solution was crystallized by adding DCM and *n*-hexane and then refrigerated for two days. Yield: 0.7 g, 92.7%; <sup>1</sup>H} NMR (CDCl<sub>3</sub>, 298 K, δ ppm): 1.3 (m, [6H], –CH<sub>3</sub>), 3.08 (m, [4H], –CH<sub>2</sub>CH<sub>3</sub>), 4.82 (m, [4H], –CH<sub>2</sub>); <sup>31</sup>P{<sup>1</sup>H} (CDCl<sub>3</sub>, 298 K, δ ppm): 87.10(s); IR (KBr) (cm<sup>-1</sup>): 1180(s), ν(P–O–C); 788(s), ν(P–O–C); 665(s), ν(P=S); 547(s), ν(P–S).

### 2.3 *In situ* solvothermal deposition of Sb<sub>2</sub>S<sub>3</sub> films from [Sb{S<sub>2</sub>P{O(Pr)}<sub>2</sub>}<sub>3</sub>]

A dispersion of [Sb{S<sub>2</sub>P{O(Pr)}<sub>2</sub>}<sub>3</sub>] (1 g) in 20 mL of ethylene glycol was prepared with continuous stirring for 1 h. Fluorine-doped tin oxide (FTO) was cleaned in an ultrasonicator using deionized water, isopropanol, and acetone prior to film deposition. The aforementioned solution was then shifted to a 25 ml Teflon-lined stainless steel autoclave, in which FTO, a conducting substrate, was submerged horizontally. The autoclave was sealed and maintained at 180 °C for 24 hours. Furthermore, the autoclave was allowed to cool down to room temperature. Subsequently, the film was removed and vacuum-dried for 24 hours. A black film was acquired. Fig. 1 depicts the design of the *in situ* solvothermal process for fabricating Sb<sub>2</sub>S<sub>3</sub> films.

### 2.4 Sb<sub>2</sub>S<sub>3</sub> film characterization

A Rigaku SmartLab diffractometer (Cu-K) was used for X-ray diffraction (XRD). The samples were scanned between 10° and 80°, with 40 kV and 50 mA supplied as the voltage and current, respectively. The generated pattern was phase-corresponded to card number 042-1393 (stibnite) in the Inorganic Crystal Structure Database (ICDD) using the PDF-2 2021 software package. OXFORD Instruments INCA Wave was used for energy-dispersive X-ray (EDX) spectroscopy and scanning electron microscopy (SEM) investigation. A Gemini/ZEISS ULTRA 55 was used for SEM, with a working distance of 5–8 mm between a sample and the pole piece as well as an acceleration voltage of





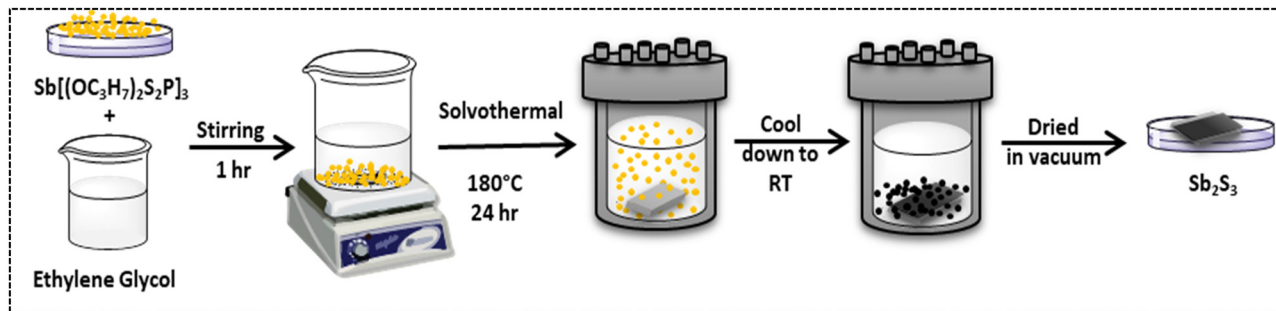


Fig. 1 Schematic representation of deposition of the  $\text{Sb}_2\text{S}_3$  film by the *in situ* solvothermal process.

5 keV. EDX was implemented with INCA-Point & ID Software and a 20 keV acceleration voltage. Raman spectroscopy was performed with the help of a Horiba Scientific Instruments along with a microscope and a 633 nm laser diode. Spectra were obtained by applying 25% laser power. The Multipurpose Analytical S/TEM JEM-F200 “F2” from Japan was operated at 200 kV to provide images for high-resolution transmission electron microscopy (HR-TEM), and selected area diffraction (SAED) patterns were obtained. A Shimadzu, UV-2600i, Japan spectrophotometer was used for the optical characterization of the  $\text{Sb}_2\text{S}_3$  films.

### 2.5 Fabrication of the $\text{Sb}_2\text{S}_3$ electrical device and characterization

As-deposited  $\text{Sb}_2\text{S}_3$  serves as a resistive switching layer in the device. A silver (Ag) top with a thickness of 200 nm was patterned on the surface of the  $\text{Sb}_2\text{S}_3$  layer by thermal evaporation. The bottom and top contacts were the FTO substrate and Ag electrode, respectively. A probe attached to a Keysight (B1500) system was used for assessing the electrical properties at room temperature and ambient pressure. Throughout all measurements, the top contact received the voltage while the bottom contact was connected to ground.

## 3. Results and discussion

### 3.1 Precursor preparation and spectroscopic studies

$[\text{Et}_3\text{NH}][(\text{OC}_3\text{H}_7)_2\text{S}_2\text{P}]$ . All characteristic bands are present in their expected regions in the IR spectrum of  $[\text{Et}_3\text{NH}][(\text{OC}_3\text{H}_7)_2\text{S}_2\text{P}]$  (Fig. S1, ESI<sup>†</sup>). All the relevant peaks were present in the  $\{^1\text{H}\}$  NMR spectrum as shown in Fig. S2 (ESI<sup>†</sup>). The  $^{31}\text{P}\{^1\text{H}\}$  NMR spectrum (Fig. S3, ESI<sup>†</sup>) exhibits a singlet at 110.66 ppm, demonstrating the symmetric character of the species in solution as well as the equivalent nature of the phosphorus nuclei.

$[\text{Sb}\{\text{S}_2\text{P}\{\text{O}(\text{Pr})_2\}_3]$ . Two strong intensity bands attributed to  $\nu(\text{P}-\text{O}-\text{C})$  and  $\nu\{\text{P}-\text{O}-(\text{C})\}$  stretching vibrations were observed in the ranges 1185–1100 and 825–790  $\text{cm}^{-1}$ , respectively. Two strong bands due to  $\nu(\text{P}=\text{S})$  or  $\nu(\text{P}-\text{S})_{\text{asym}}$  and  $\nu(\text{P}-\text{S})$  or  $\nu(\text{P}-\text{S})_{\text{sym}}$  stretching vibrations were shifted to a lower wavenumber by 30–40  $\text{cm}^{-1}$  upon comparison with the  $[\text{Et}_3\text{NH}][(\text{OC}_3\text{H}_7)_2\text{S}_2\text{P}]$ , which is consistent with the coordination of the ligand to the central Sb-atom (Fig. S4, ESI<sup>†</sup>).

The  $\{^1\text{H}\}$  NMR spectrum for the  $[\text{Sb}\{\text{S}_2\text{P}\{\text{O}(\text{Pr})_2\}_3]$  compound (Fig. S5, ESI<sup>†</sup>) was compared with that of the corresponding  $[\text{Et}_3\text{NH}][(\text{OC}_3\text{H}_7)_2\text{S}_2\text{P}]$ . In the complex, the  $-\text{CH}_3$  perceived as multiplets in the region of 1.3–4.82 ppm, display coordination shift of approximately 0.4 ppm associated with those in the corresponding salts. The  $^{31}\text{P}\{^1\text{H}\}$  NMR spectrum of the complex demonstrates a sharp singlet at 87 ppm indicating dithiophosphate group coordination to the antimony centre in the molecule<sup>43–45</sup> (Fig. S6, ESI<sup>†</sup>). This revealed the equivalent nature of phosphorous nuclei and in solution, there existed just one form of phosphorus. The  $^{31}\text{P}\{^1\text{H}\}$  NMR chemical shift for this complex was 23 ppm upfield from that of the equivalent  $[\text{Et}_3\text{NH}][(\text{OC}_3\text{H}_7)_2\text{S}_2\text{P}]$ , which represents the anisobidentate nature<sup>46</sup> of dithiophosphate moieties. Glidewell concluded that signals in the range of 82–101 ppm represent complexes with bidentate ligands whereas monodentate ligands were found in those with signals smaller than 82 ppm after evaluating the proton decoupled  $^{31}\text{P}\{^1\text{H}\}$  NMR chemical shifts of a variety of metal dialkyl dithiophosphate.<sup>47</sup> This shows that bidentate coordination of the dithiophosphate with antimony is rather robust in solution.

### 3.2 Deposition of $\text{Sb}_2\text{S}_3$ films via the *in situ* solvothermal process

Fig. 1 depicts the schematic representation of the experimental setup. Here, the *in situ* solvothermal process serve as an alternative low temperature solution-processed method. The deposition of  $\text{Sb}_2\text{S}_3$  by immersing FTO in solution of  $[\text{Sb}\{\text{S}_2\text{P}\{\text{O}(\text{Pr})_2\}_3]$  and ethylene glycol allow formation of the film under a low temperature thermal process and further avoiding any additional step of annealing. The compatibility of the precursor with an appropriate solvent to avoid further oxidation is essential, hence the choice of a solvent plays the key role in addition to variation in the various deposition parameters. In hydrothermal or solvothermal synthesis, ethylene glycol is frequently used as a solvent.<sup>48</sup> It is both an organic biocompatible solvent and a hydrophilic liquid that is water soluble. The deposited  $\text{Sb}_2\text{S}_3$  film is pure, uniform and well-adhered without the addition of any binding agent. The optimization of the deposition parameter, for example, reaction time (12 h, 18 h and 24 h) was investigated at a lower temperature (180 °C). It was observed that,  $\text{Sb}_2\text{S}_3$  films deposited at low reaction times, 12 h and 18 h, ensued the phosphorous impurity;



while the film that was deposited at 24 h enhanced the quality of the film yielding pure  $\text{Sb}_2\text{S}_3$ . After the reaction, the black film was obtained at a lower temperature (180 °C).

### 3.3 $\text{Sb}_2\text{S}_3$ film characterization

The deposited black  $\text{Sb}_2\text{S}_3$  film is reproducible and highly stable in air and moisture. The data were successfully interpreted using several characterisation techniques such as XRD, Raman spectroscopy, SEM, and UV-Vis spectroscopy.

**3.3.1 Structural studies.** XRD reveals the qualitative properties of the phase composition and crystal structure. Fig. S7 (ESI†) displays the XRD spectrum of  $\text{Sb}_2\text{S}_3$  deposited at various reaction times (12, 18, and 24 h). There was no specific change observed in the peak position of the spectrum recorded at different intervals of time (12, 18, and 24 h). Fig. 2 presents the XRD spectrum of the as-deposited  $\text{Sb}_2\text{S}_3$  with 'hkl' planes (miller indices) indicating the growth of the orthorhombic  $\text{Sb}_2\text{S}_3$  (stibnite) phase (ICDD: 42-1393)<sup>49</sup> using filtered  $\text{CuK}\alpha$  radiation (1.5406 Å). The lattice parameters of the structure were  $a = 11.24$  Å,  $b = 11.31$  Å, and  $c = 3.83$  Å. It also displays  $P_{bnm}$  space group symmetry, with the largest peaks at  $2\theta = 15.76^\circ$ ,  $17.6^\circ$ ,  $24.9^\circ$ ,  $28.62^\circ$ ,  $29.28^\circ$ ,  $32.4^\circ$ , and  $35.68^\circ$  corresponding to the (0 2 0), (1 2 0), (1 3 0), (3 2 0), (2 1 1), (2 2 1) and (2 4 0) planes, respectively. Additionally, the produced  $\text{Sb}_2\text{S}_3$  had a high crystallinity, which was shown by the strong and sharp XRD peaks. In general, antimony, sulfur, or antimony trioxide-related crystalline impurities could not be found. Furthermore, no additional  $\text{Sb}_2\text{S}_2\text{O}$  phase-specific peaks were detected, despite  $\text{Sb}_2\text{S}_3$  made at high temperatures occasionally showing these peaks.<sup>50</sup>

The Debye–Scherrer equation (eqn (1)) was applied to XRD data to determine the crystallite diameter ( $D_c$ ) of  $\text{Sb}_2\text{S}_3$ , which was found to be 43.62 nm.

$$D_c = K\lambda/\beta \cos \theta \quad (1)$$

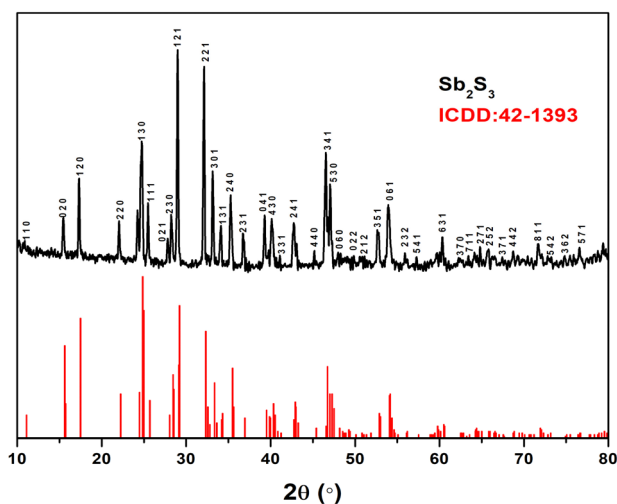


Fig. 2 XRD spectrum of the as-deposited  $\text{Sb}_2\text{S}_3$  on FTO using  $\text{Sb}[(\text{OC}_3\text{H}_7)_2\text{S}_2\text{P}]_3$  as SSP.

where  $\beta$  is the measured diffraction line's width at its half-maximum intensity,  $K$  is the shape factor, which frequently takes a value of around 0.9, and  $\lambda$  is the wavelength of the X-ray source employed in XRD.

Fig. 3 demonstrates the Raman spectrum of as-deposited  $\text{Sb}_2\text{S}_3$  on FTO. The clearly defined peaks show that the material is crystalline. It was found that the  $\text{Sb}_2\text{S}_3$  stibnite phase exhibited well-resolved Raman bands at 101, 116, 152, 192, 240, 283, and  $311 \text{ cm}^{-1}$ . The obtained peak positions correspond to the theoretically predicted  $A_g$  (at 101, 116, 152, 283, and  $311 \text{ cm}^{-1}$ ),  $B_{2g}$  (at  $192 \text{ cm}^{-1}$ ), and  $B_{1g}/B_{3g}$  (at  $240 \text{ cm}^{-1}$ ) optical modes.<sup>51,52</sup> The peaks are well-resolved, representing the crystalline nature of the sample. Well-determined Raman bands were observed at 159, 191, 240, 282, and  $311 \text{ cm}^{-1}$ , conforming to the stibnite phase. The peaks at 283 and  $311 \text{ cm}^{-1}$  arise because of the symmetric vibration of the  $\text{Sb}_2\text{S}_3$  pyramidal units having  $C_{3v}$  symmetry<sup>53,54</sup> and the peaks positioned at 192 and  $240 \text{ cm}^{-1}$  indicated the crystalline nature of  $\text{Sb}_2\text{S}_3$ .<sup>55,56</sup>

**3.3.2 Morphological studies.** The morphology of the as-deposited  $\text{Sb}_2\text{S}_3$  was explored with the help of SEM analysis. Fig. 4a presents the SEM micrograph of  $\text{Sb}_2\text{S}_3$  deposited at a reaction time of 24 h, which showed fairly bunches of rods tied together at the middle or at the end tending toward a sheaf-like morphology. Moreover, the top view of these structures displayed a hedgehog-shaped morphology. Similar morphology was observed at reaction times of 12 h and 18 h (Fig. S8a and S9a, ESI†). However, the size of the sheafs increased as the time of reaction decreased that is for 12 h ( $\sim 20 \mu\text{m}$ ), 18 h ( $\sim 40 \mu\text{m}$ ) and 24 h ( $\sim 50 \mu\text{m}$ ).

The elemental composition was ascertained by the EDX measurements. The EDX spectrum of the synthesized  $\text{Sb}_2\text{S}_3$  material confirmed the pure nature of the sample. Fig. 4c illustrates the  $\text{Sb}_2\text{S}_3$  matching EDX spectra. The samples included O, Sb, and S, as determined from the elemental composition of  $\text{Sb}_2\text{S}_3$ . Long exposure to air during analysis<sup>57</sup> or the FTO substrate may have caused the oxygen atom to exist. Moreover, according to an EDX examination, the stoichiometric ratio of Sb:S was calculated to be 2.1:3. Furthermore, the presence of phosphorus impurities decreased with an increase in the reaction time as displayed in Fig. S8b and S9b (ESI†). Fig. 4b depicts an appropriate elemental mapping of Sb and S using energy-dispersive X-ray spectroscopy (EDS), demonstrating that Sb and S were uniformly distributed across the whole  $\text{Sb}_2\text{S}_3$ . Therefore, the reaction time of 24 h is considered as the optimized time and further characterization and resistive switching were performed. Cross-sectional SEM revealed that the majority of the particles are consistently aligned with their flat sides perpendicular to the substrate surface and that the film has a thickness of between 10–20  $\mu\text{m}$  (Fig. S11, ESI†).

Fig. 5 depicts the TEM, SAED and HRTEM micrographs of the  $\text{Sb}_2\text{S}_3$  powder which were obtained after scraping from the deposited  $\text{Sb}_2\text{S}_3$  film *via* the *in situ* solvothermal process. To learn more about the shape and structure of the  $\text{Sb}_2\text{S}_3$ , additional investigation was performed using TEM, SAED, and HRTEM. Fig. S10 (ESI†) presents the size distribution



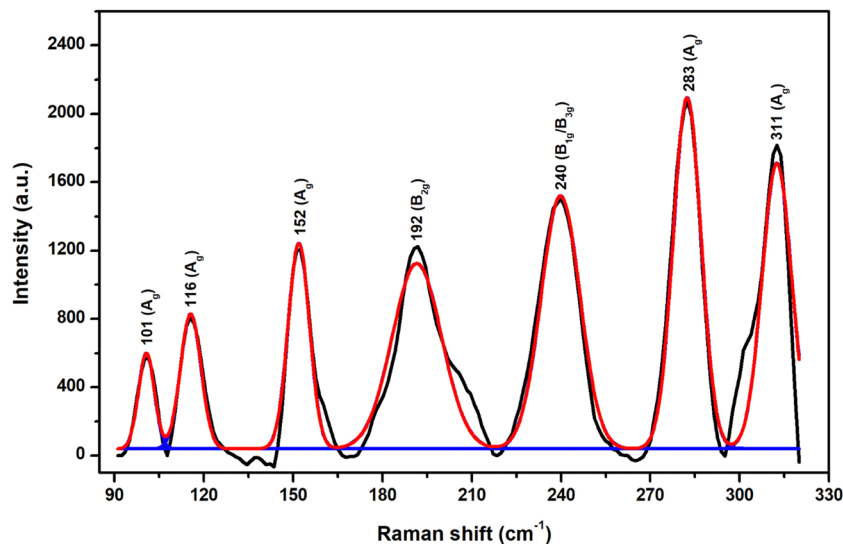


Fig. 3 Raman spectrum of the deposited  $\text{Sb}_2\text{S}_3$  on FTO using  $\text{Sb}[(\text{OC}_3\text{H}_7)_2\text{S}_2\text{P}]_3$  as SSP.

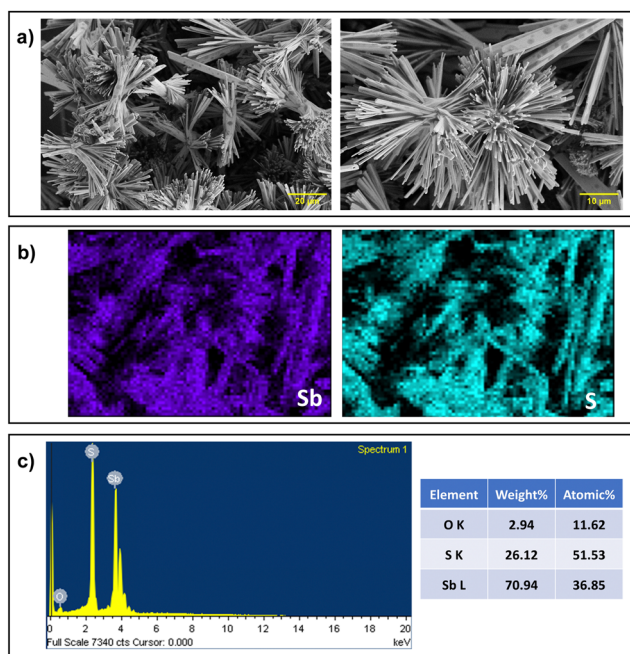


Fig. 4 (a) SEM micrograph (b) EDX elemental mapping (c) EDX spectrum with percentage of elements of the deposited  $\text{Sb}_2\text{S}_3$  film on FTO.

histogram of  $\text{Sb}_2\text{S}_3$ , which showed a mean size of the rods of 523 nm. Fig. 5a displays single crystal  $\text{Sb}_2\text{S}_3$  with a width of 467 nm. Fig. 5b and c indicates a HRTEM image of single crystal  $\text{Sb}_2\text{S}_3$ . Fig. 5b depicts the SAED pattern which indicates the single crystal  $\text{Sb}_2\text{S}_3$ . The SAED pattern of the  $\text{Sb}_2\text{S}_3$  unveils a spot design from the (120), (220), (200), and (240) planes, indicating that the  $\text{Sb}_2\text{S}_3$  have an orthorhombic phase of  $\text{Sb}_2\text{S}_3$  (ICDD:42-1393), and it resulted in a single crystal. The HRTEM image (Fig. 5c) indicates the lattice spacing of 0.36 nm corresponding to the (1 3 0) plane of orthorhombic phase of  $\text{Sb}_2\text{S}_3$ .

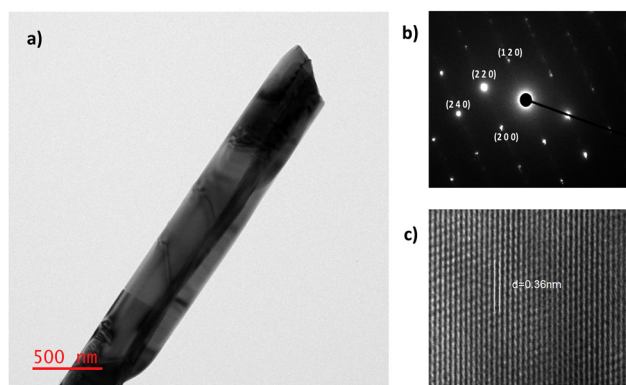


Fig. 5 (a) TEM micrograph (b) SAED pattern, and (c) phase contrast HRTEM micrograph of  $\text{Sb}_2\text{S}_3$ .

(ICDD: 42-1393). This is perpendicular to the (0 1 0) plane which indicates the direction for the growth of  $\text{Sb}_2\text{S}_3$  rods.<sup>58,59</sup>

**3.3.3 Optical studies.** UV-Vis spectroscopy in the range of 200–800 nm has been utilized to investigate the absorption of  $\text{Sb}_2\text{S}_3$ . The absorption spectrum of  $\text{Sb}_2\text{S}_3$  presented in Fig. 6a exhibits two small bands with peaks around 348 nm and 396 nm and a broad peak in the range 500–700 nm. The band gap ( $E_g$ ) energies were ascertained from the Tauc relationship as shown in eqn (2)

$$(\alpha h\nu) = A(h\nu - E_g)^n \quad (2)$$

where  $E_g$  is the bandgap energy,  $h\nu$  is the light energy,  $A$  is a constant,  $\alpha$  is the absorption coefficient, and  $n$  denotes the transition nature which equals 2 for direct transition.<sup>60</sup> When the linear portion of the curve was extrapolated from the spectra, 1.72 eV for  $\text{Sb}_2\text{S}_3$  was obtained as the band gap energy value ( $E_g$ ), as indicated in Fig. 6b. Because the morphology of the  $\text{Sb}_2\text{S}_3$  remained consistent throughout the reaction, there was no discernible alteration in the band gap. The variability in



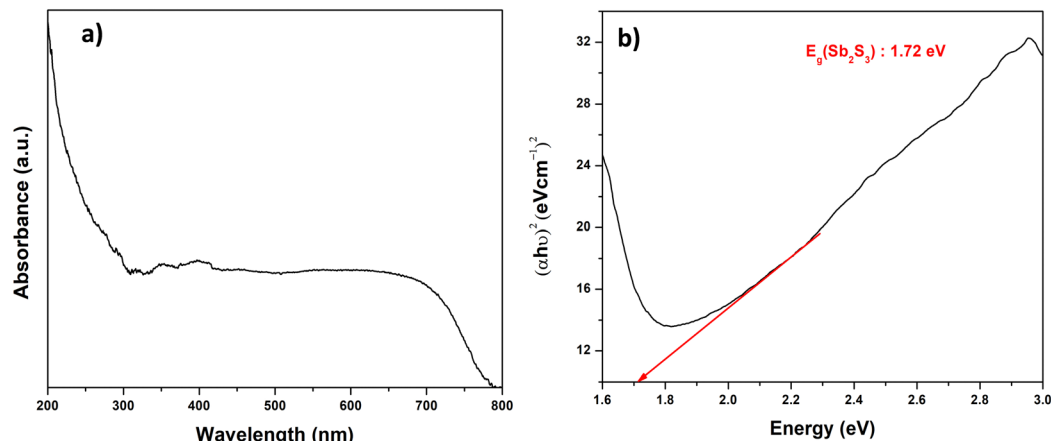


Fig. 6 (a) UV-Vis spectrum and the (b) corresponding Tauc plot of the deposited  $\text{Sb}_2\text{S}_3$  on FTO.

the crystallinity and size of the  $\text{Sb}_2\text{S}_3$  may be responsible for the variance in the band gap of the  $\text{Sb}_2\text{S}_3$ .<sup>38</sup> The band gap energies of  $\text{Sb}_2\text{S}_3$ , according to Boughalmi *et al.*, may differ depending on the technique of deposition;<sup>25</sup> meanwhile, Almalki *et al.* revealed the change in band gap energies of  $\text{Sb}_2\text{S}_3$  depending on the various solvents.<sup>58</sup>  $\text{Sb}_2\text{S}_3$  is a possible material for resistive memory switching applications since the computed band gap value was in the red portion of the solar spectrum.

### 3.4 $\text{Sb}_2\text{S}_3$ films for resistive random-access memory applications

The memristive switching properties of the as-deposited  $\text{Sb}_2\text{S}_3$  film were investigated for application in RRAM devices. Fig. 7a illustrates the structure of our memory where the  $\text{Sb}_2\text{S}_3$  film is sandwiched between the thermally evaporated silver top electrodes and the FTO bottom electrodes. The memristor was then subjected to a series of DC sweeps between the two electrodes for switching. As shown in Fig. 7b, the pristine device presents a high-resistance state (HRS) and can be switched to a low-resistance state (LRS) by applying a negative DC sweep where a SET voltage of approximately  $-1.3$  V was observed. The device can then be reset back to the HRS by applying a successive positive sweep voltage (from 0 V to 2 V). Interestingly, the subsequent switching cycle highly resembles the first one with similar SET and RESET voltages. This suggests that no extra energy is required to initialize our device and it possesses electroforming-free properties. Consecutive cyclic  $I$ - $V$  measurements were performed on the device to investigate the switching repeatability. Fig. 7c presents 80 cyclic  $I$ - $V$  curves where the memristor maintains an ON/OFF ratio of *ca.* 10 (shown in Fig. 7d). The retention of the memristor is plotted in Fig. 7e where both HRS and LRS remain stable for over 1000 s, confirming its non-volatile switching behaviour. We further investigated the device-to-device variation of our memristor. Fig. 7f plots the representative DC switching characteristic of three different  $\text{Sb}_2\text{S}_3$  memristors in this work. It can be observed that all three devices demonstrate very similar bipolar switching behaviour with negative SET and positive RESET voltages. There are some variations in the switching currents

which are likely due to the non-uniformity of the  $\text{Sb}_2\text{S}_3$  film deposited *via* our novel precursor and deposition approach. This can be improved with further optimisation of the deposition process.

The conduction mechanism of our memristor is investigated by fitting the  $I$ - $V$  curve on a log scale as shown in Fig. 7g. According to the fitting, the conduction at the HRS is dominated by the Space-Charge-Limited-Current (SCLC).<sup>1</sup> The device exhibits a linear relationship of current with applied voltage initially between 0 V and 0.4 V. This may be attributed to an Ohmic conduction mechanism resulting from thermally produced charge carriers.<sup>61</sup> The slope increases to around 2 at higher applied voltages ( $0.4 \leq V \leq 1.1$  V), and the current reveals the voltage square dependency, which is caused by the trap-controlled space charge limited current (TC-SCLC). Ohmic conduction dominates the conduction mechanism in the LRS, with the current linearly varying with the applied voltage.

To further elucidate the switching mechanism of our  $\text{Sb}_2\text{S}_3$  memristor, we conducted an area dependence investigation. Fig. 7h presents the  $I$ - $V$  characteristics for devices with areas of  $150 \mu\text{m}$  by  $150 \mu\text{m}$  and  $250 \mu\text{m}$  by  $250 \mu\text{m}$ . Despite showing similar  $I$ - $V$  curves, some discrepancies can be observed between the two devices at the OFF state. We further plot the obtained HRS and LRS as a function of device area in Fig. 7i. It is evident that the HRS decreases with the increasing device area while the LRS is independent of the area. Such area-independence in the LRS suggests the filamentary switching mechanism of our  $\text{Sb}_2\text{S}_3$  based memristor. However, the formation of the conductive filament is unlikely due to the top Ag electrode as it will require a positive bias to switch the device ON, which is in contrary to our observation. Instead, atoms originated in the as-deposited  $\text{Sb}_2\text{S}_3$  film are likely to be responsible for the resistive switching. As a cation-like species in alloys, Sb can ionize and move upon application of a sufficient electric field.<sup>21</sup> Application of a negative potential to the top electrode attracts the cationic Sb species to the top electrode, where neutral Sb atoms are accumulated and start to form a filament. This is reasonable for our memristor as our  $\text{Sb}_2\text{S}_3$  film is slightly Sb-rich (Sb:S ratio 2.1:3). The readily





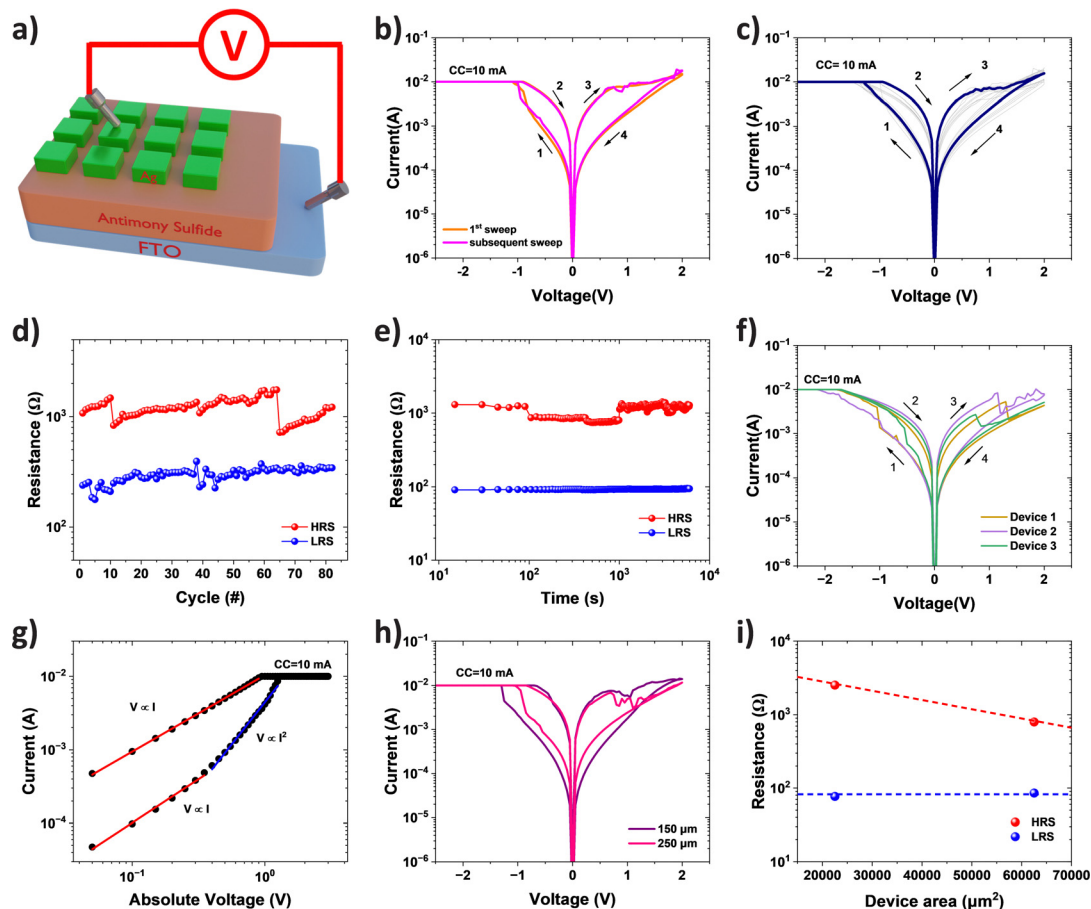


Fig. 7 Resistive switching properties of the deposited  $\text{Sb}_2\text{S}_3$  RRAM devices. (a) Schematic structure of  $\text{Sb}_2\text{S}_3$ -based memory devices. (b) The DC  $I$ - $V$  curves of the electro-forming and subsequent switching cycles. (c) 80 consecutive DC  $I$ - $V$  cycles of the device. (d) DC endurance, and (e) retention of the memristor. (f) Representative DC  $I$ - $V$  cycles from three different devices. (g)  $I$ - $V$  curve fitting using a SCLC model. (h)  $I$ - $V$  switching characteristics of the memristor with a top electrode side length of 250  $\mu\text{m}$  and 150  $\mu\text{m}$ . (i) Both HRS and LRS of the memristor as a function of device area.

available Sb atoms in the film also facilitate the initial switching process without the requirement of a larger electroforming voltage. Similar switching behaviour was also observed in other  $\text{Sb}_2\text{S}_3$  based memory devices.<sup>34</sup> We further confirmed

this mechanism by replacing Ag with W (an inert metal) as the top electrode. Similar  $I$ - $V$  characteristics were observed as shown in Fig. S12 (ESI<sup>†</sup>). The pristine memristor can be switched to a low resistance state under a similar negative bias

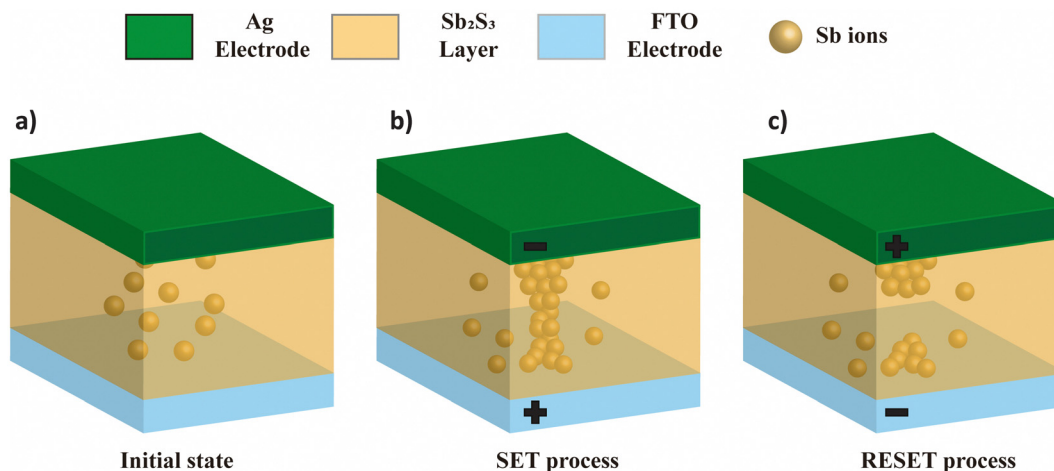


Fig. 8 Schematic of the filamentary-based resistive switching mechanism of our Ag/ $\text{Sb}_2\text{S}_3$ /FTO resistive memory cell.





Table 1 Performance comparison between various metal sulfide-based RRAM devices

S. no.	Device structure	Electroforming required	Retention (s)	On/off ratio	$V_{\text{set}}$ (V)	$V_{\text{reset}}$ (V)	Ref.
1	Ag/Sb <sub>2</sub> S <sub>3</sub> /FTO	No	10 <sup>4</sup>	10	-1.3	2	This work
2	Cu/ZnS/Pt	Yes	—	10–10 <sup>8</sup>	~0.6	~ -0.7	62
3	Cu/ZnS/ITO	Yes	—	10–10 <sup>8</sup>	~0.6	~ -1.2	62
4	Cu/MoS <sub>2</sub> /Ni–Mn–In on Si	Yes	10 <sup>3</sup>	6 × 10 <sup>2</sup>	1.36	0.98	63
5	Graphene/MoS <sub>2-x</sub> O <sub>x</sub> /graphene	Yes	~10 <sup>5</sup>	10 <sup>2</sup>	3.5	-4.8	64

and back to a high resistance state with a positive bias. This similar switching behaviour with an inert electrode further supports our argument that Sb-filament based switching is the dominant mechanism for our memristor.

Fig. 8 illustrates the schematics of the potential switching mechanism of our Sb<sub>2</sub>S<sub>3</sub> based memory device. As-deposited Sb<sub>2</sub>S<sub>3</sub> has a significant number of Sb atoms (Fig. 8a). Upon application of a strong enough electric field, these cation-like entities can ionise and move.<sup>21</sup> When a negative potential is applied, the cationic Sb species are drawn to the top electrode, where neutral Sb atoms are gathered to form the filament. When the growing filament reaches the bottom electrode, the device changes to the LRS (SET process, as illustrated in Fig. 8b). A reverse bias, on the other hand, will cause the device to return to the HRS by displacing the cationic Sb species to the bottom electrode and dissolving the prepared filament (the RESET process, as depicted in Fig. 8c).

We further compared the resistive switching performance of our Ag/Sb<sub>2</sub>S<sub>3</sub>/FTO memristor with other reported metal sulfide-based RRAM devices in Table 1. Our device has competitive performance with unique electroforming-free properties. More importantly, the successful demonstration of memristive behaviour has certainly paved the way for future large-scale manufacturing of functional chalcogenide thin films using our novel one-step, facile, low-temperature *in situ* solvothermal deposition approach.

## 4. Conclusions

In this study, we reported the *in situ* solvothermal process which serves as an alternative low temperature solution-based deposition of Sb<sub>2</sub>S<sub>3</sub> films from [Sb{S<sub>2</sub>P{O(Pr)<sub>2</sub>}<sub>3</sub>}] as a SSP to demonstrate scalable and industrially compatible processing for memristive devices. The proposed solution processible route is advantageous as it demonstrates ease of stoichiometric modulation, low equipment and energy cost and high throughput production. Here, the process was performed without involvement of any binding agent and hence the additional step of annealing is eliminated. The deposited film was pure, uniform, and well-adhered, revealing the crystalline orthorhombic stibnite phase. Significantly, the deposited Sb<sub>2</sub>S<sub>3</sub> film was employed as a novel resistive random-access memory device and it demonstrated typical bipolar resistive switching behaviour with electroforming-free properties. The Ag/Sb<sub>2</sub>S<sub>3</sub>/FTO memristive device showed reliable stability over 80 cycles while maintaining an ON/OFF ratio of *ca.* 10. The switching behaviour is attributed to the formation and rupture of Sb

ion-based filaments within the Sb<sub>2</sub>S<sub>3</sub> matrix. This further opens up wide range of possibilities for future low power and high performance devices.

## Author contributions

Sayali Shrishail Harke: conceptualization, methodology, investigation, validation, formal analysis, and writing – original draft. Tongjun Zhang: methodology and writing – review & editing. Ruomeng Huang: methodology, and writing – review & editing. Chitra Gurnani: conceptualization, methodology, validation, formal analysis, writing – review & editing, and supervision.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The authors are grateful to Mahindra University, Hyderabad, for providing the necessary funds and experimental resources to conduct the work. The authors thank SAIF Chandigarh Punjab University for access to NMR characterization facility. We also acknowledge the School of Physics and School of Chemistry, University of Hyderabad, for TEM and SEM characterisation, respectively.

## References

- A. H. Jaafar, L. Meng, Y. J. Noori, W. Zhang, Y. Han, R. Beanland, D. C. Smith, G. Reid, K. De Groot, R. Huang and P. N. Bartlett, *J. Phys. Chem. C*, 2021, **125**, 26247–26255.
- R. Huang, K. Sun, K. S. Kiang, K. A. Morgan and C. H. De Groot, *Microelectron. Eng.*, 2016, **161**, 7–12.
- A. H. Jaafar, M. O'Neill, S. M. Kelly, E. Verrelli and N. T. Kemp, *Adv. Electron. Mater.*, 2019, **5**, 1900197.
- M. J. Lee, C. B. Lee, D. Lee, S. R. Lee, M. Chang, J. H. Hur, Y. B. Kim, C. J. Kim, D. H. Seo, S. Seo, U. I. Chung, I. K. Yoo and K. Kim, *Nat. Mater.*, 2011, **10**, 625–630.
- D. I. Son, D. H. Park, J. Bin Kim, J.-W. Choi, T. W. Kim, B. Angadi, Y. Yi and W. K. Choi, *J. Phys. Chem. C*, 2011, **115**, 2341–2348.
- G. U. Kamble, A. P. Patil, R. K. Kamat, J. H. Kim and T. D. Dongale, *ACS Appl. Electron. Mater.*, 2023, **5**, 2454–2481.



- 7 V. K. Nagareddy, M. D. Barnes, F. Zipoli, K. T. Lai, A. M. Alexeev, M. F. Craciun and C. David Wright, *ACS Nano*, 2017, **11**, 3010–3021.
- 8 P. Priyadarshini, S. Das and R. Naik, *RSC Adv.*, 2022, **12**, 9599–9620.
- 9 V. Gupta, S. Kapur, S. Saurabh and A. Grover, *IETE Tech. Rev.*, 2020, **37**, 377–390.
- 10 K. C. Kwon, J. H. Baek, K. Hong, S. Y. Kim and H. W. Jang, *Nano-Micro Lett.*, 2022, **14**, 1–30.
- 11 W. Banerjee, *Electronics*, 2020, **9**, 1–24.
- 12 Z. Jiang, Y. Wu, S. Yu, L. Yang, K. Song, Z. Karim and H. S. P. Wong, *IEEE Trans. Electron Devices*, 2016, **63**, 1884–1892.
- 13 K. H. Nam, J. H. Kim, W. J. Cho, C. H. Kim and H. B. Chung, *J. Nanosci. Nanotechnol.*, 2016, **16**, 10393–10396.
- 14 J. H. Lee, S. P. Park, K. Park and H. J. Kim, *Adv. Funct. Mater.*, 2020, **30**, 1907437.
- 15 E. J. Yoo, M. Lyu, J. H. Yun, C. J. Kang, Y. J. Choi and L. Wang, *Adv. Mater.*, 2015, **27**, 6170–6175.
- 16 A. P. Ranavavare, S. J. Kadam, S. V. Prabhu, S. S. Chavan, P. V. Anbhule and T. D. Dongale, *Mater. Lett.*, 2018, **232**, 99–102.
- 17 C. Schindler, M. Meier, R. Waser and M. N. Kozicki, *Proc. – 2007 Non-Volatile Mem. Technol. Symp. NVMTS 07*, 2007, pp. 82–85.
- 18 N. I. Mou and M. Tabib-Azar, *Appl. Surf. Sci.*, 2015, **340**, 138–142.
- 19 H. Hönigschmid, M. Angerbauer, S. Dietrich, M. Dimitrova, D. Gogl, C. Liaw, M. Markert, R. Symanczyk, L. Altimime, S. Bournat and G. Müller, *IEEE Symp. VLSI Circuits, Dig. Tech. Pap.*, 2006, vol. 42, pp. 110–111.
- 20 N. Banno, T. Sakamoto, T. Hasegawa, K. Terabe and M. Aono, *Jpn. J. Appl. Phys., Part 1*, 2006, **45**, 3666–3668.
- 21 S. Yoo, T. Eom, T. Gwon and C. S. Hwang, *Nanoscale*, 2015, **7**, 6340–6347.
- 22 T. D. Dongale, N. D. Desai, K. V. Khot, N. B. Mullani, P. S. Pawar, R. S. Tikke, V. B. Patil, P. P. Waifalkar, P. B. Patil, R. K. Kamat, P. S. Patil and P. N. Bhosale, *J. Solid State Electrochem.*, 2017, **21**, 2753–2757.
- 23 P. S. Pawar, R. S. Tikke, V. B. Patil, N. B. Mullani, P. P. Waifalkar, K. V. Khot, A. M. Teli, A. D. Sheikh and T. D. Dongale, *Mater. Sci. Semicond. Process.*, 2017, **71**, 102–108.
- 24 L. Zhang, C. Wu, W. Liu, S. Yang, M. Wang, T. Chen and C. Zhu, *J. Mater. Chem. A*, 2018, **6**, 21320–21326.
- 25 R. Boughalmi, A. Boukhachem, M. Kahlaoui, H. Maghraoui and M. Amlouk, *Mater. Sci. Semicond. Process.*, 2014, **26**, 593–602.
- 26 P. Makreski, G. Petruševski, S. Ugarković and G. Jovanovski, *Vib. Spectrosc.*, 2013, **68**, 177–182.
- 27 K. F. Abd-El-Rahman and A. A. A. Darwish, *Curr. Appl. Phys.*, 2011, **11**, 1265–1268.
- 28 F. Aousgi and M. Kanzari, *Energy Procedia*, 2011, **10**, 313–322.
- 29 X. Wang, R. Tang, C. Wu, C. Zhu and T. Chen, *J. Energy Chem.*, 2018, **27**, 713–721.
- 30 P. Zhong, J. Xie, R. Bagheri, Q. Yi, Q. Chen, J. Tan, L. He, F. Zhang, L. Zhang and G. Zou, *Chem. Commun.*, 2019, **55**, 14530–14533.
- 31 O. Savadogo and K. C. Mandal, *Sol. Energy Mater. Sol. Cells*, 1992, **26**, 117–136.
- 32 W. Yi, F. Perner, M. S. Qureshi, H. Abdalla, M. D. Pickett, J. J. Yang, M. X. M. Zhang, G. Medeiros-Ribeiro and R. S. Williams, *Appl. Phys. A: Mater. Sci. Process.*, 2011, **102**, 973–982.
- 33 W. Wu, B. Shan, K. Feng and H. Nan, *Mater. Sci. Semicond. Process.*, 2016, **44**, 18–22.
- 34 A. G. Wallace, R. P. King, N. Zhelev, A. H. Jaafar, W. Levason, R. Huang, G. Reid and P. N. Bartlett, *Electrochim. Acta*, 2022, **432**, 141162.
- 35 C. Gurnani, S. L. Hawken, A. L. Hector, R. Huang, M. Jura, W. Levason, J. Perkins, G. Reid and G. B. G. Stenning, *Dalton Trans.*, 2018, **47**, 2628–2637.
- 36 M. A. Malik, N. Revaprasadu and P. O'Brien, *Chem. Mater.*, 2001, **13**, 913–920.
- 37 J. Rodriguez-Castro, P. Dale, M. F. Mahon, K. C. Molloy and L. M. Peter, *Chem. Mater.*, 2007, **19**, 3219–3226.
- 38 G. Murtaza, M. Akhtar, M. Azad Malik, P. O'Brien and N. Revaprasadu, *Mater. Sci. Semicond. Process.*, 2015, **40**, 643–649.
- 39 D. C. Onwudiwe, O. C. Olatunde and S. Mathur, *Phys. B*, 2021, **605**, 412691.
- 40 W. Lou, M. Chen, X. Wang and W. Liu, *Chem. Mater.*, 2007, **19**, 872–878.
- 41 J. R. Castro, K. C. Molloy, Y. Liu, C. S. Lai, Z. Dong, T. J. White and E. R. T. Tiekink, *J. Mater. Chem.*, 2008, **18**, 5399–5405.
- 42 A. L. Bingham, J. E. Drake, C. Gurnani, M. B. Hursthouse, M. E. Light, M. Nirwan and R. Ratnani, *J. Chem. Crystallogr.*, 2006, **36**, 627–630.
- 43 J. L. Lefferts, K. C. Molloy, J. J. Zuckerman, I. Haiduc, C. Guta and D. Ruse, *Inorg. Chem.*, 1980, **19**, 1662–1670.
- 44 I. Haiduc, D. B. Sowerby and S. F. Lu, *Polyhedron*, 1995, **14**, 3389–3472.
- 45 M. Wieber and M. Schröpf, *Phosphorus, Sulfur Silicon Relat. Elem.*, 1995, **102**, 265–277.
- 46 J. S. Yadav, R. K. Mehrotra and G. Srivastava, *Polyhedron*, 1987, **6**, 1687–1693.
- 47 C. Glidewell, *Inorg. Chim. Acta*, 1977, **25**, 159–163.
- 48 Y. J. Zhu and F. Chen, *Chem. Rev.*, 2014, **114**, 6462–6555.
- 49 P. G. Sheikhiabadi, M. Salavati-Niasari and F. Davar, *Mater. Lett.*, 2012, **71**, 168–171.
- 50 W. D. Xiang, G. Y. Ji, W. Jing, Y. X. Yang and X. N. Liu, *Glass Phys. Chem.*, 2012, **38**, 245–253.
- 51 M. Delaney, I. Zeimpekis, D. Lawson, D. W. Hewak and O. L. Muskens, *Adv. Funct. Mater.*, 2020, **30**, 1–10.
- 52 R. Parize, A. Katerski, I. Gromyko, L. Rapenne, H. Roussel, E. Kärber, E. Appert, M. Krunks and V. Consonni, *J. Phys. Chem. C*, 2017, **121**, 9672–9680.
- 53 R. G. Sotelo Marquina, T. G. Sanchez, N. R. Mathews and X. Mathew, *Mater. Res. Bull.*, 2017, **90**, 285–294.
- 54 J. S. Eensalu, K. Tõnsuaadu, I. Oja Acik and M. Krunks, *Mater. Sci. Semicond. Process.*, 2022, **137**, 106209.



- 55 B. B. H. Juárez, S. Rubio, J. Sánchez-dehesa and C. Lopez, *Adv. Mater.*, 2002, **14**, 1486–1490.
- 56 S. M. Hwang, J. Kim, Y. Kim and Y. Kim, *J. Mater. Chem. A*, 2016, **4**, 17946–17951.
- 57 S. Mourdikoudis and L. M. Liz-Marza, *Chem. Mater.*, 2013, **25**, 1465–1476.
- 58 R. Almalki, E. M. Mkawi and Y. Al-Hadeethi, *J. Mater. Sci.: Mater. Electron.*, 2020, **31**, 9203–9211.
- 59 G. Shen, D. Chen, K. Tang, X. Jiang and Y. Qian, *J. Cryst. Growth*, 2003, **252**, 350–354.
- 60 M. Lajnef and H. Ezzaouia, *Open Appl. Phys. J.*, 2009, **2**, 23–26.
- 61 R. Huang, X. Yan, K. A. Morgan, M. D. B. Charlton and C. H. De Groot, *J. Phys. D: Appl. Phys.*, 2017, **50**, 175101.
- 62 F. Zhuge, K. Li, B. Fu, H. Zhang, J. Li, H. Chen, L. Liang, J. Gao, H. Cao, Z. Liu and H. Luo, *AIP Adv.*, 2015, **5**, 1–8.
- 63 A. Kumar, S. Pawar, S. Sharma and D. Kaur, *Appl. Phys. Lett.*, 2018, **112**, 1–5.
- 64 M. Wang, S. Cai, C. Pan, C. Wang, X. Lian, Y. Zhuo, K. Xu, T. Cao, X. Pan, B. Wang, S. J. Liang, J. J. Yang, P. Wang and F. Miao, *Nat. Electron.*, 2018, **1**, 130–136.

