

# Phase change properties of ternary AgSbSe<sub>2</sub> chalcogenide films

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Despite a low transition temperature the ternary AgSbSe<sub>2</sub> alloy possesses phase change properties including larger resistance change, a single crystalline structure and smaller volume change upon crystallization. Basic functionality of a reversible switching by a pulsed laser has been demonstrated.

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## 1. Introduction

Amorphous chalcogenide films are the most attractive materials for the wide application of micro-optical, micro-mechanical and optical/electronic memories *etc* [1-6]. Among them, tellurium-based alloys are regarded as a class of suitable materials for memory applications [7-8]. Recently, remarkable high electrical resistance change upon transition was found in binary selenium-based alloys such as In-Se or Sb-Se systems [9-11]. Here, we report the electrical and structural properties as well as a reversible switching test in ternary AgSbSe<sub>2</sub> films. The results show this ternary selenium-based compound possesses a desired performance and might act as a novel class of alternative for the application in the optical/electronic memory.

## 2. Experimental

Thin films of AgSbSe<sub>2</sub> were fabricated on glass or Si substrates by evaporating single element sources in an MBE chamber [12]. Transition temperatures were determined by temperature dependent sheet resistance measurements in a protecting Ar-atmosphere. All X-ray-measurements were performed at room temperature (Philips X'Pert MRD System). Density and thickness changes upon crystallization were obtained from XRR. Local switching trial in the AgSbSe<sub>2</sub> films was performed using a static tester ( $\lambda = 830$  nm) with an objective lens of high numerical aperture (N.A. = 0.8). Reflectance change of the irradiated region was monitored and the topography of the irradiated region was subsequently identified using an atomic force microscope (Dimension 3100, DI).

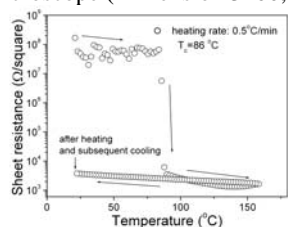


Fig. 1. Temperature-dependent sheet resistance measurement of a 72 nm AgSbSe<sub>2</sub> film on a glass substrate.

Fig. 1 shows the temperature-dependent sheet resistance of a 72 nm AgSbSe<sub>2</sub> film on a glass substrate taken at a heating rate of 0.5 °C/min. A sharp drop of the sheet resistance of more than four orders of magnitude is observed at 86 °C, which is later identified as an amorphous-to-crystalline transition by X-ray diffraction (XRD). The considerable noise at low temperature in the curve comes from the extremely high resistance value of the amorphous film, which is beyond the limit ( $\sim 10^8$  Ω/square) of the setup.

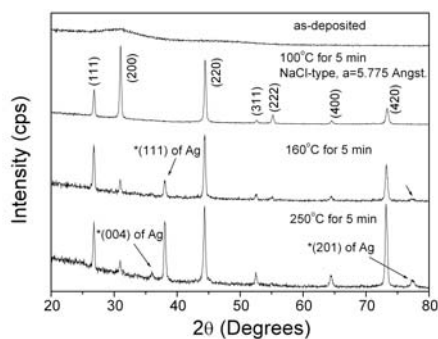


Fig. 2. XRD-scans of a 100 nm AgSbSe<sub>2</sub> film on a glass substrate in the as-deposited state and after annealing at different temperatures.

Fig. 2 depicts the XRD-scans of a 100 nm AgSbSe<sub>2</sub> film on a glass substrate in the as-deposited state and after annealing at different temperatures. The as-deposited films are confirmed to be in the amorphous state while a single crystalline phase with a NaCl-type structure is formed after annealing at 100 °C. The lattice parameter of the NaCl-type structure is determined to be  $a = 5.775 \pm 0.005$  Å, which agrees nicely with the values of 5.786 Å previously reported for single-crystalline AgSbSe<sub>2</sub> [13]. The XRD-data also confirm that the sharp drop at 86 °C in Fig. 1 corresponds to the amorphous-to-crystalline transition. Further annealing at temperatures higher than 160 °C will cause phase separation, which was identified

to be a segregation of crystalline silver out from the NaCl-typed matrix.

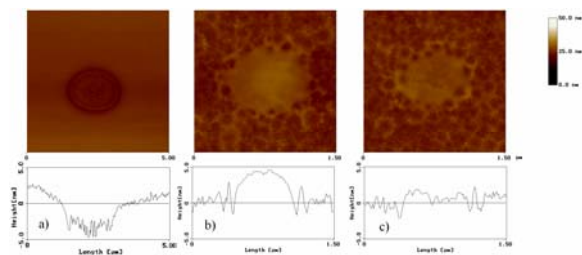


Fig. 3. a) An AFM image of a crystalline bit in an 87 nm as-deposited AgSbSe<sub>2</sub> film produced by laser of 15 mW and 80  $\mu$ s, and its typical cross section. b) Amorphous bit written with a laser pulse of 20 mW and 150 ns into a crystalline surrounding. c) Perfect erasure by a subsequent laser pulse of 4 mW and 80 ns.

### 3. Phase change recording

A reversible recording/erasing trial is performed by a pulsed laser and AFM, as described elsewhere [14-17]. Fig. 3a shows an AFM image and a typical cross section of a crystalline mark. The crystalline bit is produced by a laser pulse of 15 mW and 80  $\mu$ s. Crystallization does not only lead to a reflectivity increase but also an increase in film density [18]. Therefore the crystallized region corresponds to a depression in the amorphous surrounding. The modified area in Fig. 3a has a circular shape with a diameter of around 1.7  $\mu$ m. The saturation depth of the bit is  $4.2 \pm 0.5$  nm and the thickness of the as-deposited amorphous film is  $87 \pm 3$  nm. This leads to a change in thickness of  $4.8 \pm 0.8\%$ , which is in good agreement with the density change of 3.8% determined by XRR [12]. Fig. 3b is a micrograph of an amorphous bit produced by the write pulse in the crystalline matrix. The amorphous bit is created by the laser of 20 mW and 150 ns. Besides a decrease in reflectance, amorphization leads also to a decrease in density and therefore to a local increase in film thickness [18]. The diameter of the amorphous bit is around 650 nm and the height is approximately 4.4 nm. Fig. 3c shows an image of an erased bit produced by a subsequent pulse of 4 mW and 80 ns. After the irradiation, the height of the erased bit has already recovered to the same height as the crystalline background, indicating that complete erasure is realized.

### 4. Conclusions

In summary, despite a low transition temperature the ternary AgSbSe<sub>2</sub> alloy possesses desired phase change properties including larger resistance change, a single crystalline structure and smaller volume change upon crystallization. Basic functionality of a reversible switching by a pulsed laser is successfully demonstrated, indicating that selenide might play an active role in the application of phase change memory.

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