

Non-volatile memory device based on AgI-Ag₃BO₃ thin film

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ABSTRACT

This work employs AgI-Ag₃BO₃ to fabricate nonvolatile memory device. First the AgI-Ag₃BO₃ amorphous material is prepared with quick cooling method. The thin film is grown onto Pt electrode with pulsed laser deposition. The resulting memory device presents a superiorly high difference between the low and high resistivity states, with a ratio above 10⁶. In addition, the device shows a high stability after more than 10 thousand cycles. This structure may be used as a memory device with fast response and high stability.

Keywords - nonvolatile memory, pulsed laser deposition, solid electrolyte, AgI-Ag₃BO₃

I. INTRODUCTION

The rapid development of scale-down in the new era has advanced plenty of novel paradigms of designs, e.g., high performance electrical devices [1]. A straightforward method to work at small scale is to make the constituent materials smaller [2, 3]. Another approach is to find nanoscaled materials that as the elemental building blocks [4, 5]. Other tactics involve the infrastructures of multiple distinctly disparate ingredients to achieve functional properties at the macroscopic level [6-9]. The electrical equipment and electronics are experiencing revolutions in different fields [10-12]. For example, memories are needed almost everywhere in our daily life and work space, such as smart phones, computers, and servers. Due to the explosion of data, the demand for high performance and density memories are continuously increasing, approaching the final limit of Moore's law. In the last few decades, many endeavors have been put into the novel nonvolatile memory devices. Solid electrolyte materials attracted quite a lot of attention due to their unique characteristics that are superior or comparable with tradition materials [13-16]. A unique characteristic of solid electrolyte is that its resistivity may alter within a huge range, possibly 7 orders of magnitude. These distinct states of resistivity can be used as the „1“ and „0“ in memory devices. The successfully generated

devices have shown superior performance compared with traditional equipment [17, 18].

In this work, a novel AgI-Ag₃BO₃ solid electrolyte system was generated to fabricate nonvolatile device. To the best of authors' knowledge, this system has not been applied before.

II. EXPERIMENTS

AgI, AgNO₃, and H₃BO₃ are thoroughly mixed with molar ratio 1:3:1 and ground into fine powders at room temperature (RT). The mixture is heated to 900 °C and kept for 2 hours. The chemical reaction is:



The mixture is then quenched to RT into a bulk target with a chemical composition of 1:1 AgI-Ag₃BO₃.

Pulsed laser deposition was performed on the bulk target at a vacuum below 10⁻⁴ Pa. The laser fluence is 220 mJ with a frequency of 5 Hz. The total deposition time is 7 minutes.

III. RESULTS

3.1. Characterization

The XRD pattern (Fig. 1) shows that the generated material is apparently amorphous with a very small content of crystalline fractions. The characteristic peak with a circle is α-AgI, while the square is Ag(111). Among the two common phases of AgI, β-AgI has a lower conductivity than α-AgI. There have been a few reports on the applications of α-AgI, e.g., Tatsumisago et al. prepared nanocrystalline α-AgI in glassy networks at RT [19].

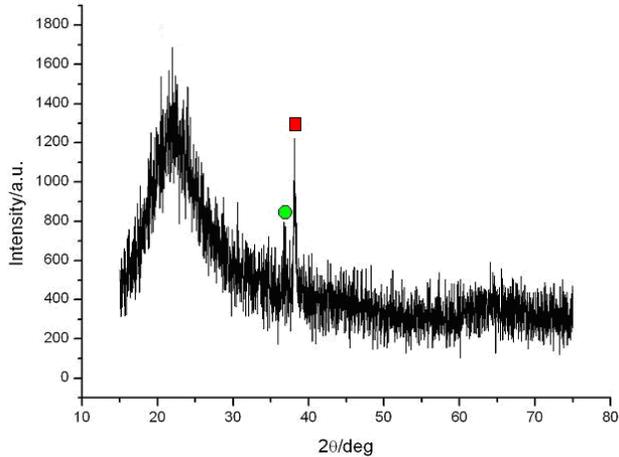


Fig. 1. XRPD pattern of the generated AgI-Ag₃BO₃

The generated thin film was examined under TEM and the image is presented in Fig. 2. Based on the image, the AIB thin film shows a morphology of nanocrystallites.

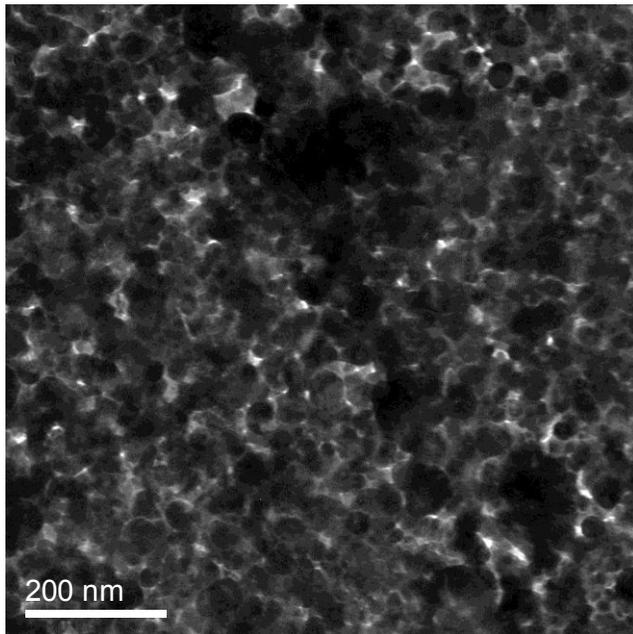


Fig. 2. TEM image of the AgI-Ag₃BO₃ thin film

The X-ray photoelectron spectroscopy (XPS) of the AgI-Ag₃BO₃ thin film is shown in Fig. 3. Based on the results, the Ag atom in the thin film is close to metal, indicating that it has a weak interaction with I. As a result, it may be easily converted to free Ag in electrical field.

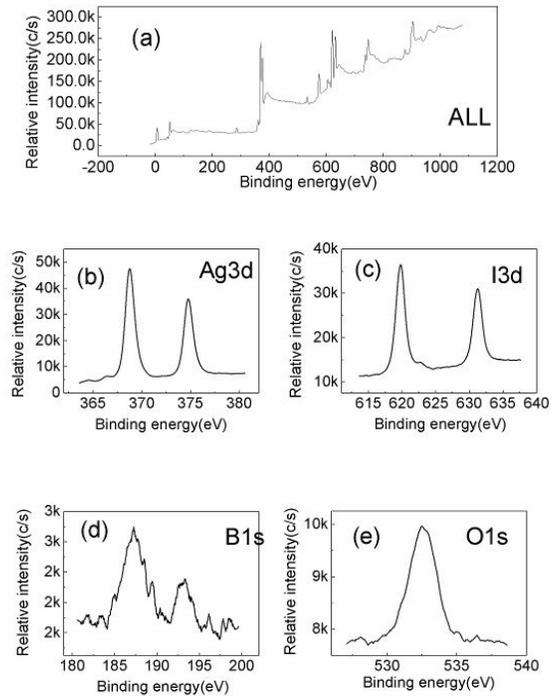


Fig. 3. XPS results of the AgI-Ag₃BO₃ thin film

Fig. 4 shows the structure of the final memory device based on AgI-Ag₃BO₃. The substrate at the bottom electrode is Pt and the top electrode is Ag. The thickness of Ag is about 100 nm and the AIB thin film is ~250 nm. The nonconductive SiO₂ was deposited on the Pt electrode and drilled with focused ion beam etching.

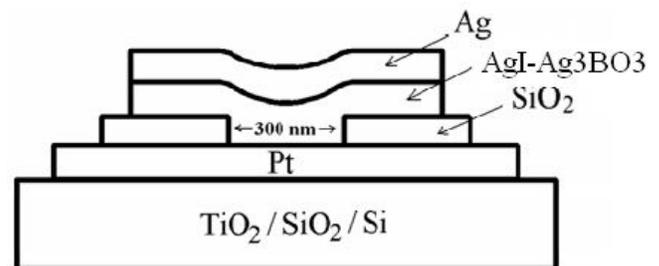


Fig. 4. Structure of the memory device based on AgI-Ag₃BO₃

The on/off switching effect of the memory device based on AgI-Ag₃BO₃ thin film is shown in Fig. 5. The ratio between high and low resistivity is above 10⁶ and the device is cycled for more than 10⁴ times.

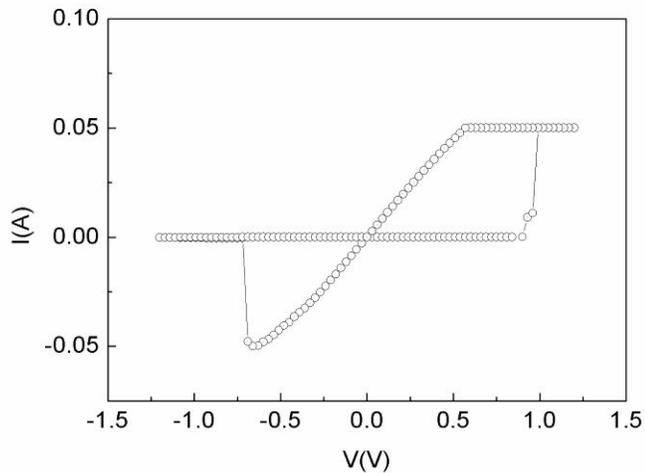


Fig. 5. The on/off switching of the memory device based on AIB thin film

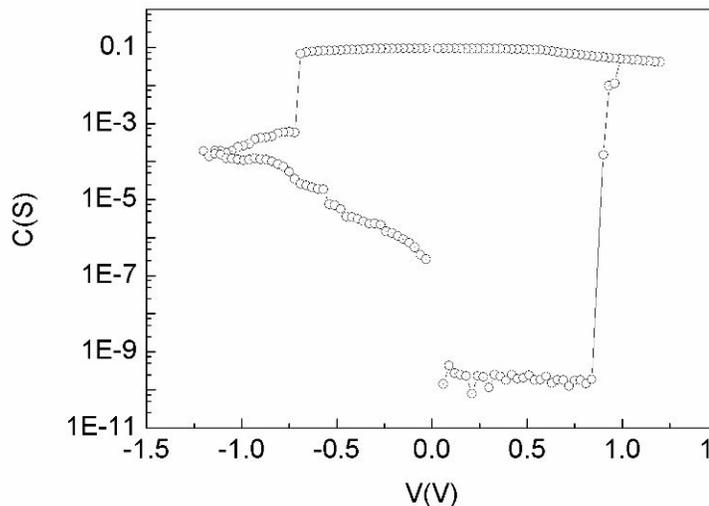


Fig. 6. Conductivity of the device under different voltages

Fig. 7 presents the on/off characteristics at the read-write pulse of 1 kHz. In each cycle, a positive voltage of 2.5 V as read pulse is applied to the device to enter the state of low resistivity; a negative voltage of 8 V as erase is applied to switch the device to the high resistivity state. Two pulses of read at 2 V are designated after write and erase. Between any two pulses, the voltage is 0. All pulses have a width of 0.125 ms and one full cycle is 1 ms. Based on the current which is 0 after read pulse, the state is high resistivity; while after write pulse, the current is approximately 11 mA, indicating the low resistivity state.

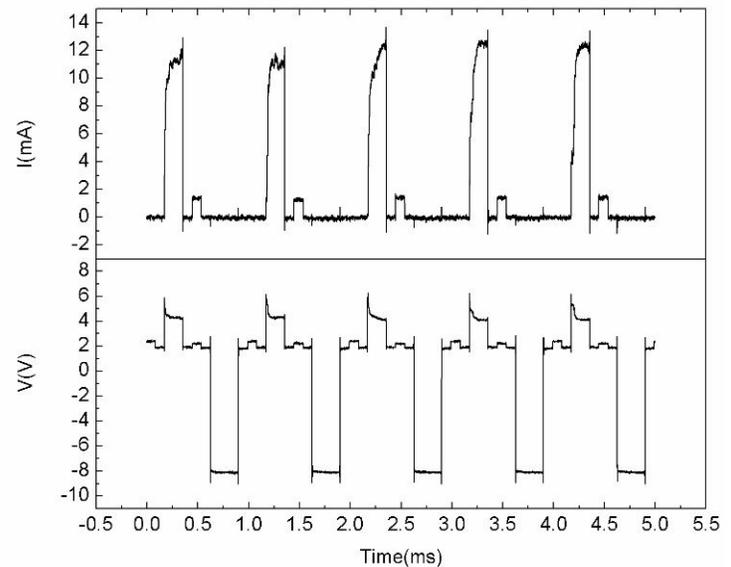


Fig. 7. AIB thin film device read-write cycles under 1 kHz

IV. CONCLUSION

This study prepares AgI-Ag₃BO₃ amorphous solid electrolyte via the quick cooling method. Then the AgI-Ag₃BO₃ thin film is grown onto Pt electrode with pulsed laser deposition. The resulting memory device shows a great difference between the low and high resistivity states, with a ratio above 10⁶. In addition, the device shows a high stability after more than 10 thousand cycles. This infrastructured device may be used as a memory device with fast response and high stability.

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