ZrN-Based Flexible Resistive Switching Memory

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Abstract — In this letter, ZrN-based resistive random access memory (RRAM) is investigated for flexible memory applications for the near future. Due to the room-temperature fabrication process, the device is suitable for low-temperature flexible and mobile technologies. The TiN/ZrN/TiN device exhibits excellent AC endurance cycling (10^4), a rapid speed (45 ns) and stable retention (10^4 s) at 100°C without any degradation. In addition, RRAM devices built with an additional HfN interface layer exhibit small operational voltage variations and stable switching characteristics. The flexibility of the device is excellent, and it maintains excellent electrical characteristics at a bending radius of up to 4 mm.

Index Terms — RRAM, nitrogen ions, conductive filament, resistive switching.

I. INTRODUCTION

Resistive random access memory (RRAM) has received a substantial amount of attention for use in high-density memory applications [1, 2]. Flexible electronics are becoming an evolving technology due to their attractive properties and high degree of flexibility for portable electronic devices that can be used in daily life. Flexible electronic devices are fabricated on plastic substrates and not Si substrates, so they are low-cost and green electronic devices [3]. It is well known that high local temperature and self-heating effects in the filament region can result in uncontrollable filament issues, such as switching current overshoot and reliability problems [4]. Various transition metal nitrides, such as HfN, ZrN, VN, SiN, SiCN, and CrN, have been used as either resistive switching (RS) layers or interface layers in RRAM devices [2, 5-8]. Transition metal nitride ZrN has a high thermal conductivity (50 W/mK) and a semiconducting phase [8]. These properties are beneficial for suppressing local heating effects and improving the controllability of the conductive filament. Several types of flexible RRAM have been reported but are not appealing due to their poor endurance, high bending radius, low speeds and poor bending cycling behavior [9-17]. Kim et al. [9] reported flexible RRAM, which showed good retention (10^4 s) but poor endurance (150 cycles) and a high bending radius (10 mm). Qian et al. [10] also reported flexible devices with poor endurance (800 cycles), poor retention (10^3 s), a high bending radius (14 mm), and poor bending cycling behavior (850 cycles). To overcome these problems in flexible devices, we propose a flexible RRAM based on a new TiN/HfN/ZrN/TiN structure that has shown excellent characteristics, such as AC endurance (10^7 cycles), speed (45 ns), bending radius (4 mm) and retention (10^4 s) at 100°C. These characteristics indicate that the TiN/HfN/ZrN/TiN RRAM stack is better than the previously reported RRAM stacks for flexible nonvolatile memory applications. Table 1 shows a comparison of oxide-based flexible devices with current work.

In this work, we demonstrate a new TiN/ZrN/TiN device on a polyethylene terephalate (PET) substrate for emerging non-volatile RRAM for flexible applications. The highly flexible TiN/ZrN/TiN device shows excellent switching characteristics with an on/off resistance ratio (>10^5), device speed (45 ns), AC endurance (10^7 cycles) and stable retention (10^4 s) at 100°C. The RS mechanism of the device is based on the formation and rupture of conductive filaments (CFs) in the ZrN layer due to nitrogen vacancies. In addition, we used HfN as a thin layer that is inserted between the top electrode and ZrN layer to reduce the voltage variation in the device.

II. DEVICE FABRICATION

First, the TiN bottom electrode (BE) was deposited by sputtering on a flexible PET substrate. A 10 nm-thick ZrN RS layer was deposited on the TiN BE by rf sputtering. Finally, a 100 nm TiN top electrode (TE) with a diameter of 100 μm using a metal shadow mask was deposited by sputtering at room temperature to form a TiN/ZrN/TiN device. In addition, 3 nm HfN was used as an interface layer between the TiN and ZrN for comparison to suppress the operational voltage variation. An Agilent B1500A semiconductor parameter analyzer was used for the DC measurements, and an AC pulse was generated by an Agilent B1530A waveform generator/fast measurement

Table 1: Comparison of previous reported oxides and organic materials based flexible devices with current work.

<table>
<thead>
<tr>
<th>Device Structure</th>
<th>Endurance (Cycles)</th>
<th>Speed</th>
<th>Retention</th>
<th>Bending radius (mm)</th>
<th>Bending cycles</th>
<th>Report Year</th>
</tr>
</thead>
<tbody>
<tr>
<td>NiSnOx/ATO/PET</td>
<td>10^3</td>
<td>10^3 s</td>
<td>10^3 s @ 85°C</td>
<td>5</td>
<td>10^3</td>
<td>2013</td>
</tr>
<tr>
<td>CuWOx/TiO</td>
<td>10^3</td>
<td>10^3 s</td>
<td>10^3 s @ 85°C</td>
<td>5,5</td>
<td>10^3</td>
<td>2016</td>
</tr>
<tr>
<td>Ag/ITO/ATO/PET</td>
<td>10^3</td>
<td>10^3 s</td>
<td>10^3 s @ 85°C</td>
<td>5</td>
<td>10^3</td>
<td>2016</td>
</tr>
<tr>
<td>ZnSnOx/AZO/AgO</td>
<td>10^4</td>
<td>10^4 s</td>
<td>10^4 s</td>
<td>10</td>
<td>10^4</td>
<td>2013</td>
</tr>
<tr>
<td>Pt/TiO2/ATO/PET</td>
<td>600</td>
<td>10^4 s</td>
<td>10^4 s</td>
<td>20</td>
<td>10^4</td>
<td>2017</td>
</tr>
<tr>
<td>TiOxSnOx/PTE</td>
<td>800</td>
<td>10^4 s</td>
<td>10^4 s</td>
<td>14</td>
<td>850</td>
<td>2017</td>
</tr>
<tr>
<td>Ni/AlOx/Ni</td>
<td>200</td>
<td>10^4 s</td>
<td>10^4 s</td>
<td>5</td>
<td>10^4</td>
<td>2018</td>
</tr>
<tr>
<td>AlPdTiO/PET</td>
<td>500</td>
<td>10 ms</td>
<td>10^4 s</td>
<td>5</td>
<td>10^4</td>
<td>2019</td>
</tr>
<tr>
<td>Al/Parylene/W</td>
<td>500</td>
<td>10 μm</td>
<td>10^4 s</td>
<td>10</td>
<td>500</td>
<td>2016</td>
</tr>
<tr>
<td>TiN/HfN/ZnN/TiN</td>
<td>10^4</td>
<td>45 ns</td>
<td>10^4 s @ 100°C</td>
<td>4 mm</td>
<td>10^4</td>
<td>This work</td>
</tr>
</tbody>
</table>

This work was supported by Ministry of Science and Technology, Taiwan, under Project No. NSC 105-2221-E-009-134 -MY3.
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III. RESULTS AND DISCUSSION

Fig. 1(a) shows the bipolar RS behavior of the TiN/ZrN/TiN device with a SET voltage (~1.3 V) and a RESET voltage (~0.8 V). Figure 1(b) depicts the voltage distribution of the device during 100 continuous DC switching cycles. The device exhibits a wide variation in the SET/RESET voltages, in the low voltage region (0 to 0.4 V) of the HRS, the thermally generated carrier migration inherent in the nitride film causes a low current, and the fitted curve shows an Ohmic conduction dependence (I-V). As the voltages of the TiN/ZrN/TiN and TiN/HfN/ZrN/TiN devices rise to (0.4 to 1.3 V) and (0.4 to 0.9 V), respectively, the number of injected carriers exceeds the number of thermally generated carriers and trapping and detrapping of the inherent defects in the nitride, which leads to the conduction following the SCLC behavior of trap filling (I ~ V²). At high voltages (>1.3 V for TiN/ZrN/TiN and >0.9 V for TiN/HfN/ZrN/TiN), all traps are filled with charge carriers, a conductive path is formed, and the current suddenly increases; the devices are SET from the HRS to the LRS. In the entire LRS region of both devices, the I-V curves are linear with a slope of 1, which is consistent with the Ohmic conduction mechanism. In standard SCLC theory, the current density J emitted by trap-controlled SCLC can be expressed as:

\[ J = (\theta / \theta + 1) (9/8) \varepsilon E (V^2 / L^3) \]

where \( \theta = (N_C / N_i) \exp(E_i / kT) \) is the ratio of the free electrons to the trapped electrons, \( N_C \) is the effective density of states in the conduction band, \( N_i \) is the number of empty electron traps, \( \varepsilon \) is the dielectric constant of free space, \( E_i \) is the relative permittivity, \( \mu \) is the electron mobility, \( V \) is the applied voltage, and \( L \) is the film thickness.

We also present a schematic model of a TiN/ZrN/TiN device based on the formation and rupture of CFS, which is shown in Figs. 2 (b) and (c). The CF consists of nitrogen (N) vacancies in the device that are created mostly at the grain boundaries of the RS layer during the forming or SET process. The creation of N vacancies in ZrN devices is very similar to the creation of oxygen vacancies in metal oxide RRAM devices [4]. After forming or the SET process, the free N ions released from ZrN remain at the TiN/ZrN interface rather than being absorbed by TiN because the Gibbs free energy of TiN (~217 kJ/mol) is higher than that of ZrN (~336 kJ/mol), and nitrogen ions need more energy to be absorbed in TiN [18]. During the RESET process, the nitrogen ions drift back to the ZrN and recombine with the nitrogen vacancies, causing the filament to rupture. Due to the high thermal conductivity (50 W/mK) of ZrN, the local heating effect is not as severe as that in the metal...
oxide device, and the shape and size of the filament can be better controlled [8].

To control the switching voltage variation and switching stability of the TiN/ZrN/TiN device, we used a 3 nm HfN thin layer between the TE and 7 nm ZrN layer. The improved characteristics of the TiN/HfN/ZrN/TiN device are shown in Fig. 3. The device has SET and RESET voltages of approximately 0.8 and −0.6 V, respectively (Fig. 3a). The TiN/HfN/ZrN/TiN device has a smaller SET/RESET voltage change than the TiN/ZrN/TiN device (Fig. 3b). For TiN/HfN/ZrN/TiN devices with a speed of 45 ns, an AC endurance of up to $10^5$ cycles is achieved (Fig. 3c) using a pulse height of 1.6 V for the SET operation and −1.5 V for the RESET operation. The retention of the TiN/HfN/ZrN/TiN device is measured at 100 °C, and both the LRS and HRS exhibit high stability (Fig. 3d).

Fig. 4 (a) and (b) illustrate the CF model for the TiN TE, N vacancies are generated in the ZrN layer and migrate to the TiN BE to form conical filaments in the ZrN layer. As the N vacancies in the ZrN layer increase, ZrN becomes electrically conductive and behaves like a "virtual electrode"[19]. It should be noted that the conductive virtual electrode, which is part of the CF, also consists of N vacancies. Now, the "virtual electrode" in the ZrN layer tends to be the rupture point and recovery locations in the matrix layer near the HfN/ZrN interface. After the device switches to the HRS, the root of the CF does not dissolve completely during the RESET process. In the TiN/HfN/ZrN/TiN device, the CF is preferably connected or ruptures at the HfN/ZrN interface, and the ZrN layer behaves as a virtual electrode [19]. This RS mechanism enhances the uniformity and reduces the SET/RESET voltage variations of the TiN/HfN/ZrN/TiN device.

The flexible properties of the TiN/HfN/ZrN/TaN device are also investigated. During the measurements, the device is bent up to surface curvature radius of 25–2 mm. Figure 4(c) shows the values for various bending radii. The device exhibits a stable resistance ratio between the LRS and HRS without any degradation to mechanical bending from 25 to 4 mm but rapid degradation occurs for 2 mm bending. This degradation in the resistances is due to the increase in the sheet resistance of the RS layer and BE during the bending condition [20].

Fig. 4 (d) shows the endurance characteristics of the device after bending from a flat state to a 4 mm radius 12,000 times. Both the LRS and HRS are highly stable up to 10,000 bending cycles, after which the device shows degradation in both states. The first read operation is performed after the first bending cycle, and then a read operation is performed after every 200 bending cycles to measure the stability of the device. The AC endurance test is also investigated for a 4 mm bending radius at a read voltage of 0.3 V, and the result is shown in Fig. 4 (e). By using a pulse height of 1.6 V for the SET process and −1.5 V for the RESET process with a pulse width of 45 ns, the flexible device can maintain both LRS and HRS for up to $10^7$ cycles without any degradation. The inset of Fig. 4(e) exhibits the consistency of LRS and HRS values among the measured results for the 20 devices. The high stability of the LRS and HRS confirms the high reproducibility of the device.

For experimental evidence, we obtain SEM images of the device in the flat and bent conditions, as shown Fig. 5. In Fig. 5 (a), there are no cracks visible for the device in the flat condition. For the device with a bending R = 4 mm, the crack density is much less, as observed from Fig. 5 (b), and the increase in the resistance of the ZrN layer due to the few cracks results in the maintenance of a good memory performance [13, 21]. In Fig. 5 (c), it is clear that there are many cracks in the device when R = 2 mm. These cracks lead to an increase in the ZrN layer resistance on the ITO/PET substrate, resulting in degradation of the device performance. Therefore, the device shows rapid degradation in the LRS and HRS when the device is bent to R = 2 mm.

IV. CONCLUSION

We investigated RRAM devices based on a ZrN RS layer. The device has excellent switching characteristics, such as a high AC durability and stable retention at high temperatures. In addition, the device fabricated with a HfN interfacial layer between the TE and ZrN exhibits small operational voltage variations and stable switching characteristics. The flexible device shows excellent switching characteristics when bent up to a radius of 4 mm. Finally, this report clearly establishes that the ZrN-based RRAM device studied herein has potential for flexible memory applications.
REFERENCES


