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Enhanced resistive switching memory characteristics and mechanism using a Ti nanolayer at the W/TaO_x interface

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Abstract

Enhanced resistive memory characteristics with 10,000 consecutive direct current switching cycles, long read pulse endurance of $>10^5$ cycles, and good data retention of $>10^4$ s with a good resistance ratio of $>10^2$ at 85°C are obtained using a Ti nanolayer to form a W/TiO_x/TaO_x/W structure under a low current operation of 80 μA, while few switching cycles are observed for W/TaO_x/W structure under a higher current compliance >300 μA. The low resistance state decreases with increasing current compliances from 10 to 100 μA, and the device could be operated at a low RESET current of 23 μA. A small device size of 150 × 150 nm^2 is observed by transmission electron microscopy. The presence of oxygen-deficient TaO_x nanofilament in a W/TiO_x/TaO_x/W structure after switching is investigated by Auger electron spectroscopy. Oxygen ion (negative charge) migration is found to lead to filament formation/rupture rather than oxygen vacancy (hole) migration, and it is controlled by Ti nanolayer at the W/TaO_x interface. Conducting nanofilament diameter is estimated to be 3 nm by a new method, indicating a high memory density of ≈100 Tbit/in^2.

Keywords: Resistive switching; W/TaO_x; Ti nanolayer; Oxygen ion migration; Nanofilament

Background

Resistive switching random access memories (RRAM) with simple metal-insulator-metal stacks are under intensive investigation owing to their great promise for use in next-generation memory applications [1-5]. However, their nonuniformity in switching, low yield, and unclear switching mechanism hinder their practical realization. RRAM devices with simple composition, easy fabrication process, and good 3D integration compatibility will be needed in the future. Methods such as doping, formation polarity control, bottom electrode modification, nanocrystal insertion, and interfacial engineering have recently been investigated to improve the characteristics of resistive switching memory [6-10]. Among other important switching materials such as TiO_x [11,12], NiO_x [13], HfO_x [10,13-15], ZrO_x [16-22], $Na_{0.5}Bi_{0.5}TiO_3$ [23], $SrTiO_3$ [24], ZnO [25,26], GeO_x [27], and SiO_x [28], tantalum oxide (TaO_x) is one of the most promising choices for future RRAM applications. However, TaO_x-based RRAM devices are infrequently reported [5,29-34]. Terai et al. [32] used a TiO_2 layer in a Ru/Ta_2O_5/TiO_2/Ru stack with good thermal stability. Ninomiya et al. [33] reported an Ir/$Ta_2O_{5-\delta}$/TaO_x/TaN structure, and Lee et al. [5] reported a Pt/Ta_2O_{5-x}/TaO_{2-x}/Pt crossbar structure with two layers of TaO_x and at least one of the inert electrodes such as Ru, Ir, and Pt. Generally, many researchers use one inert electrode to improve the performance of resistive switching memory [5,34]; however, tungsten (W) as both bottom and top electrodes in a W/TiO_x/TaO_x/W structure has not yet been reported. In this work, a resistive switching memory device using a Ti nanolayer at the W/TaO_x interface and enhanced memory characteristics such as excellent 10,000 consecutive stable dc switching cycles, long read pulse endurance of $>10^5$ cycles, and good data retention of 10^4 s at 85°C with a large resistance ratio of $>10^2$ under a low compliance current (CC) of 80 μA are reported. Furthermore, the device can be operated with a small 'RESET' current of 23 μA. For comparison, the W/TaO_x/W memory device is also fabricated. The device size of 150 × 150 nm^2 is observed using a high-resolution transmission

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electron microscope (HRTEM). The thicknesses of TiO_x and TaO_x nanolayers are 3 and 7 nm, respectively. The presence of oxygen-deficient TaO_x conducting filaments is investigated by Auger electron spectroscopy (AES) before and after switching of the memory devices. The switching mechanism of the oxygen ion migration rather than the vacancy migration due to a lower barrier height of electrons is investigated, and a filament diameter of ≈3 nm is calculated using a new method also reported in this work. Considering a small filament diameter, a high memory density of ≈100 Tbit/in.^{2} could be designed in future.

Methods

W/Ti/TaO_x/W-structured (device S1) and W/TaO_x/W-structured (device S2) resistive switching memory stacks were fabricated. A small via size of 150 × 150 nm^2 was etched into the SiO_2 on the W bottom electrode (BE), which was about 100 nm in thickness. A high-κ Ta_2O_5 film with a thickness (t_{Ta2O5}) of ≈7 nm was then deposited by an e-beam evaporator, followed by the sequential deposition of a thin (≈3 nm) interfacial layer of titanium (Ti) and ≈200-nm-thick W layer as a top electrode (TE) by rf sputtering. For device S2, no Ti layer was deposited. The final devices were obtained after a lift-off process. Memory device structure and thicknesses of all layers were observed by transmission electron microscopy (TEM) with an energy of 200 keV. The TaO_x material was also confirmed by quadrupole secondary ion mass spectroscopy (SIMS; ATOMIKA SIMS 4500, MA-Tek, Hsinchu, Taiwan) which had a high depth resolution. Primary beam was O^{2+} with an energy of 0.5 keV and analysis area of 37.5×37.5 μm^2. A bias was applied to the TE, and the BE was electrically grounded. Pristine S1 and S2 devices were electroformed by applying positive voltage to the TE before consecutive resistive switching cycle measurements.

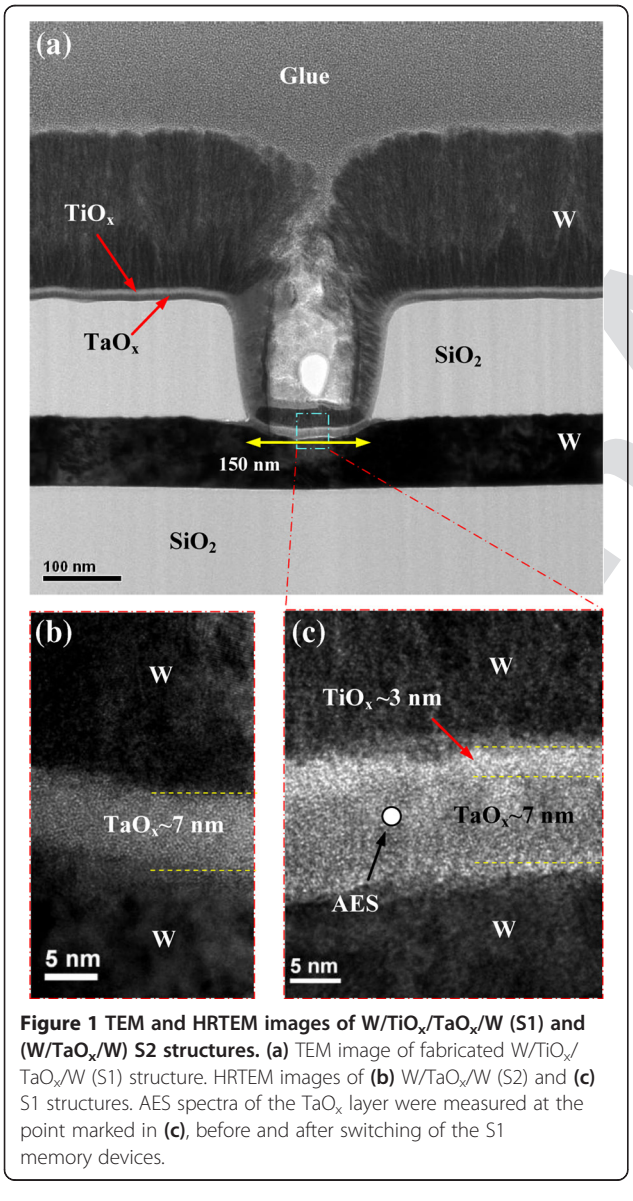

Figure 1 TEM and HRTEM images of W/TiO_x/TaO_x/W (S1) and (W/TaO_x/W) S2 structures. (a) TEM image of fabricated W/TiO_x/TaO_x/W (S1) structure. HRTEM images of **(b)** W/TaO_x/W (S2) and **(c)** S1 structures. AES spectra of the TaO_x layer were measured at the point marked in **(c)**, before and after switching of the S1 memory devices.

Results and discussion

Figure 1a shows a typical cross-sectional TEM image of the W/TiO_x/TaO_x/W structure. The device size is 150 × 150 nm^2. HRTEM images of the S1 and S2 devices are shown in Figure 1b,c. The thicknesses of the TiO_x and

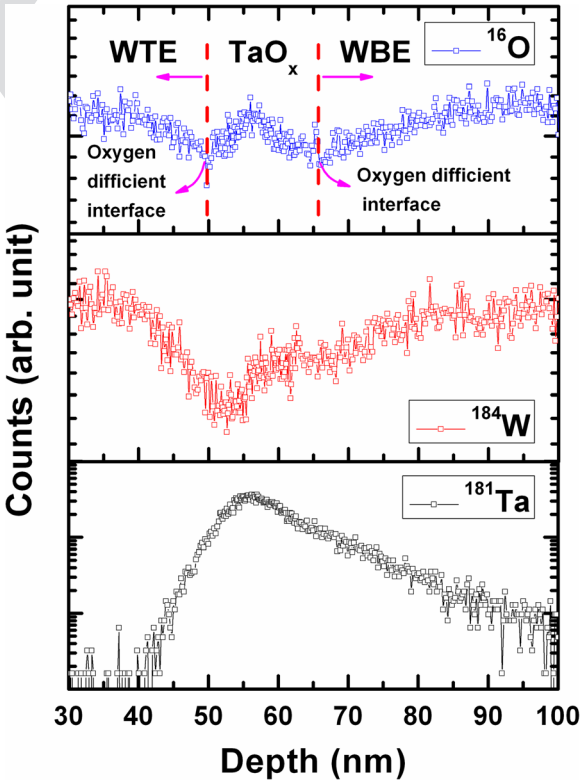

Figure 2 SIMS depth profile of W/TaO_x/W structure.

TaO_x layers are approximately 3 and 7 nm, respectively, and both films show an amorphous characteristic. Figure 2 shows typical SIMS depth profiles of ^{16}O, ^{184}W, and ^{181}Ta materials. The thickness of the TaO_x layer is about 15 nm; however, this is higher than the deposited film thickness of 7 nm. This is due to the trail effect and surface roughness of W BE, as we can see from the depth of 57 to 65 nm (or approximately 7 nm) of the ^{184}W depth profile. It is interesting to note that the TaO_x/W interface is found to be an oxygen-deficient layer, which makes it a more conducting interface. On the other hand, the conducting filament will be formed after breaking the Ta-O bonds in the bulk Ta_2O_5 layer rather than the W/TaO_x interface. This is because the Ta_2O_5 layer is more insulating than the W/TaO_x interface, so the electric field will drop across the Ta_2O_5 film rather than the W/TaO_x interface which probably results into multi-filaments or an uncontrolled nanofilament diameter. As Ti removes oxygen from the Ta_2O_5 film in the W/TiO_x/TaO_x/W structure, the film becomes more oxygen-deficient TaO_x, which is vital to achieve an improved resistive switching. Considering Gibbs free energies of TiO_2, Ta_2O_5, and WO_3 films, which are −887.6, −760.5, and −506.5 kJ/mol, respectively, at 300 K [35], Ti will consume the highest

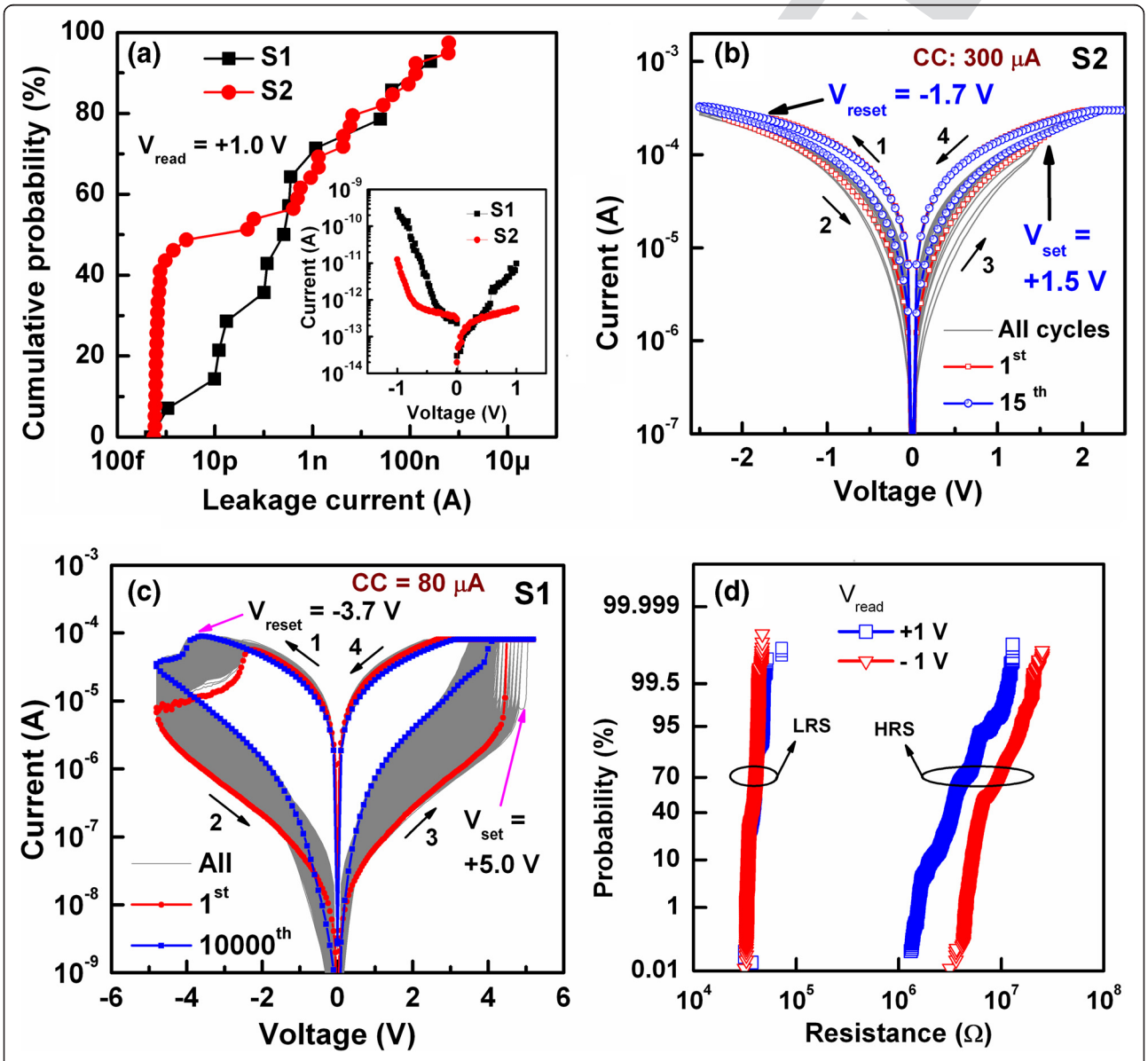

Figure 3 Cumulative probabilities of leakage currents and LRS/HRS and switching I-V curves of S1 and S2. (a) Cumulative probability of leakage currents for S1 and S2 devices with typical via size of 0.8 μm. Inset: leakage current vs. voltage characteristics. Switching I-V curves of **(b)** S2 and **(c)** S1 devices. The S2 device shows instability after a few cycles, while 10,000 consecutive switching cycles are observed for the S1 device. **(d)** Cumulative probability of LRS and HRS under V_{read} of ±1 V for the S1 devices.

oxygen content owing to its stronger reactivity than those of the other materials, thereby forming a Ta-rich (or defective TaO_x) film. This also prevents oxidation of the W TE at the TaO_x/W interface of device S1 owing to the migration of oxygen from the underlying films towards the Ti film, which contributes to the improved resistive switching memory performance as will be described.

The leakage current values of most of the S1 pristine devices at a read voltage (V_{read}) of 1 V are higher than that of the S2 devices because of the presence of more oxygen vacancies in the TaO_x layer owing to the oxygen-getter nature of the TiO_x layer (Figure 3a). Typical current-voltage (*I-V*) curves (inset of Figure 3a) of both devices were asymmetrical with higher current at a negative voltage (≈281 pA for S1 and ≈12.6 pA for S2 at V_{read} = −1 V) compared with that measured at a positive voltage (≈9.8 pA for S1 and ≈0.6 pA for S2 at V_{read} = 1 V). This suggests that the W TE/TaO_x interface has more oxygen vacancies than the TaO_x/W BE interface, owing to oxygen migration towards W TE during deposition. The ideal leakage current is plotted in Figure 4a and is explained as follows. It is reported that the work function (Φ_m) of W and bandgap (E_g) of amorphous Ta_2O_5 and TiO_2 are 4.55 [36], 4.2 [37], and 3.3 eV [38], respectively. The conduction band offsets of Ta_2O_5 and TiO_2 with Si are 0.3 [39] and 0.9 eV [40], respectively. Taking the electron affinity of Si as 4.05 eV, the electron affinities of Ta_2O_5 and TiO_2 are calculated to be 3.75 and 3.15 eV, respectively. The corresponding energy diagram is shown in Figure 4a as solid lines. Considering that the E_g of TiO_2

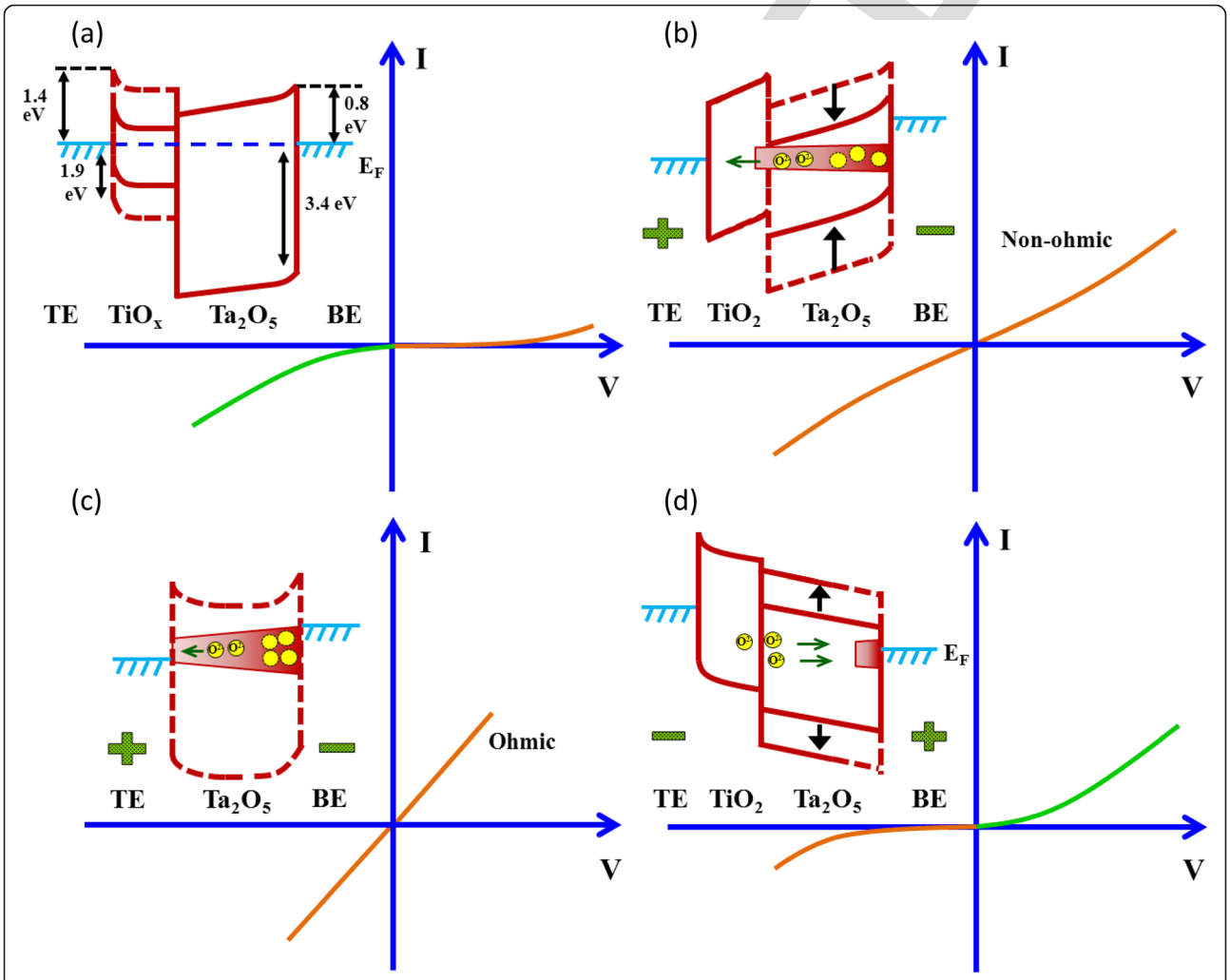

Figure 4 Schematic illustration of switching mechanism using simple energy band diagrams with I-V characteristics. (a) Energy bands of pristine S1 structure device and ideal leakage current. For TiO_2, the solid line is due to the smaller energy gap of the defective TiO_x film. I-V of LRS and corresponding energy bands with conducting filaments for **(b)** S1 and **(c)** S2 devices. For TaO_x, the solid line is due to the lower energy gap caused by the presence of oxygen-deficient filament. The S1 devices show non-ohmic I-V due to TiO_2 layer formation at the W/TaO_x interface. **(d)** Filament oxidation and leakage current at HRS are shown for the W/TiO_x/TaO_x/W devices. Filament formation/rupture is controlled by the TiO_2 layer due to O^{2-} ion migration.

for the pristine S1 device will be much lower because of oxygen vacancy creation during the deposition of W TE, the band diagram is shown in dotted lines (Figure 4a). In this case, electron injection dominates rather than hole injection because of a lower barrier height for electrons than for holes (0.8 to 1.4 vs. 3.4 eV). Both S1 and S2 devices show bipolar resistive switching behaviors. The S2 device shows few switching cycles with a higher leakage current of ≈10 μA at V_{read} = 1 V and a higher CC of 300 μA (Figure 3b). In this case, negatively charged oxygen ions (O^{2-}) migrate from the switching material towards W TE, and this has a lesser possibility to form an oxygen-rich layer at the W TE/TaO_x interface, leading to the formation of multi-conduction filaments. In the same way, no resistive switching is observed under negative forming voltage for either the S1 or S2 devices because oxygen ions migrate towards the W BE and permanent breakdown is observed (not shown here). However, the insertion of a thin (≈3 nm) Ti layer in between the W and TaO_x layers in the S1 device makes a vast difference because Ti can be used as an oxygen reservoir. Moreover, the S1 device exhibits >10,000 consecutive repeatable dc switching cycles with a better resistance ratio of 10^2 under a low CC of 80 μA (Figure 3c). However, a thicker Ti layer (5 nm) results in unstable switching cycles because it gets more oxygen and behaves as an insulating layer. This indicates that the thinner (3 nm) Ti layer will control the current overflow as well as will control the filament diameter. The yield of the S1 device is very high (>95%), while that of the S2 device is very low (approximately 10%). In addition, the S2 device cannot be switched below a CC of 300 μA and shows an ohmic behavior, while the S1 device shows switching even at a low CC of 10 μA (discussed later) with non-ohmic current conduction. The average values and standard deviation/average are found to be 39.7 and 0.11, 38.4 kΩ and 0.08 for low-resistance state (LRS) and 1.9 and 2.11, 8.6 MΩ and 0.43, for high-resistance state (HRS) at V_{read} of 1 V and −1 V, respectively (Figure 3d). This suggests that the LRS has a tighter distribution than the HRS because of the formation of the TiO_2 layer, which will have a higher E_g than the pristine one. Similarly, the leakage current at V_{read} of −1 V is lower than that at +1 V because of the lower electron injection barrier at the TE/TiO_2 interface than that at the BE/TaO_x interface after switching. Under 'SET', O^{2-} rather than oxygen vacancies will migrate from TaO_x towards the TE, resulting in a TiO_2 layer which controls the conducting vacancy filament diameter in the TaO_x layer by controlling current overflow and producing a tighter distribution of the LRS (Figure 4b). Owing to this series resistance, the S1 devices exhibit non-ohmic-simulated ideal current, as shown in Figure 4b, whereas an ohmic current is observed for the S2 devices under SET (Figure 4c). It is true that the conducting filament is formed through the TaO_x film

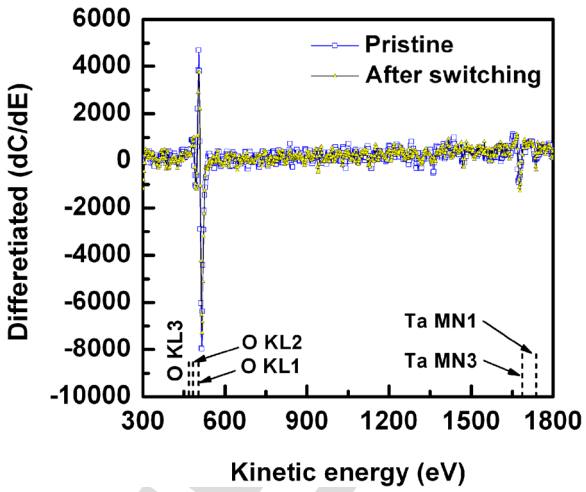

Figure 5 Differentiated (dC/dE) AES spectra vs. kinetic energy for pristine and after-switching RRAM devices. The spectra are from a typical via size of 0.4 × 0.4 μm^2 and measured inside the via regions. An oxygen-deficient (or Ta-rich) TaO_x layer is observed after few switching cycles, confirming oxygen-deficient TaO_x filament formation after SET.

(Figure 4b,c), which is also confirmed by AES spectra of the TaO_x film for pristine and after-switching S1 devices (Figure 5). The spectrum positions are also shown in Figure 1c. Ta-MN (1,737 and 1,680 eV) and O-KL (468, 483, and 503 eV) are observed, which confirms the formation of a TaO_x layer in the S1 device. The atomic percentages of Ta-MN3 and O-KL1 are 37.38% and 62.62% for the pristine device and 44.69% and 55.31% for the switched device, respectively. Basically, this decrease in oxygen content and increase in Ta content after switching is the evidence that an oxygen-deficient (or Ta-rich) filament is formed owing to oxygen ion migration

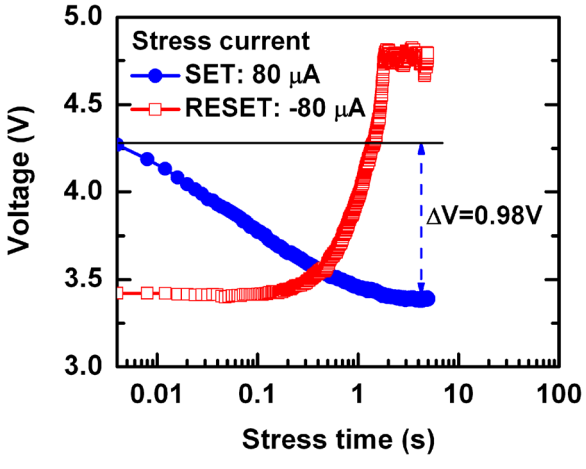

Figure 6 Voltage shift vs. stressing time under a current of ±80 μA for SET/RESET operations. The conducting filament diameter is calculated to be approximately 3 nm.

rather than oxygen vacancy migration as well as the lower energy gap of the TaO_x layer, as shown by the dotted line in Figure 4b. When negative voltage is applied to the TE, oxygen ions are pushed from the TiO_2 layer towards the conducting filament where they recombine with oxygen vacancies or oxidize the conducting filament. The device will be in HRS (Figure 4d). Control of oxygen-deficient filament formation and rupture is facilitated by insertion of the thin Ti layer at the TE/TaO_x interface, which results in repeatable and reproducible resistive switching characteristics.

The conducting filament diameter is estimated using a new method under a constant current stress of 80 μA (Figure 6). The voltage decreases (or increases) under positive (or negative) current stress after a SET (or RESET) operation. Under SET, the oxygen ions formed by the breakage of Ta-O bonds migrate towards the TiO_x layer and create an oxygen-deficient (or conducting path) in the TaO_x layer. Assuming a cylindrical nanofilament, the diameter (D) can be estimated as [41]:

$$D = \sqrt{\frac{4.q.t_{Ta_2O_5}}{\pi.\varepsilon_{Ta_2O_5}.\Delta V}}, \quad (1)$$

where ΔV (changes in the voltage shift) is found to be 0.98 V (Figure 6), q is the electronic charge (1.602×10^{-19} C), and $\varepsilon_{Ta_2O_5}$ is the dielectric permittivity of amorphous Ta_2O_5 film ($\varepsilon_{Ta_2O_5} \approx 20$ to 25). Considering all values in Equation 1, the diameter of the nanofilament is approximately 2.9 to 2.6 nm. This suggests that the present resistive switching memory device can be scaled down to <3 nm. Previously reported diameters of 5 to 10 nm for Pt/TiO_2/Pt [12], ≈15 nm for Ti/Fe:$SrTiO_3$/Nb:$SrTiO_3$ [37], and ≈1,000 nm for Pt/CuO/Pt [42] are slightly closer and higher than our calculated values, likely owing to the use of different structures as well as materials. Further study may be needed to clearly understand these results. Figure 7a shows the resistive switching characteristics with different CCs from 10 to 100 μA. The low-resistance state decreases with increasing CCs from 10 to 100 μA (Figure 7a,b), which will be useful for multi-level data storage applications. As the filament diameter increases with higher CCs, the low-resistance state decreases, and the value of RESET voltage increases. The RESET current can be scaled down to 23 μA at a low CC of 10 μA, which will be useful to a low-power operation RRAM in the near future. Our novel device also has a long read pulse endurance of $>10^5$ cycles and excellent data retention of $>10^4$ s with a good resistance ratio of $>10^2$ at 85°C at a low CC of 80 μA, as shown in Figure 8. A data retention of $>10^3$ s is also observed for a low CC of 10 μA (not shown here). Considering the obtained nanofilament diameter of approximately 3 nm, a high-density (≈100 Tbit/in.^{2}) nanoscale nonvolatile memory can be achievable in future.

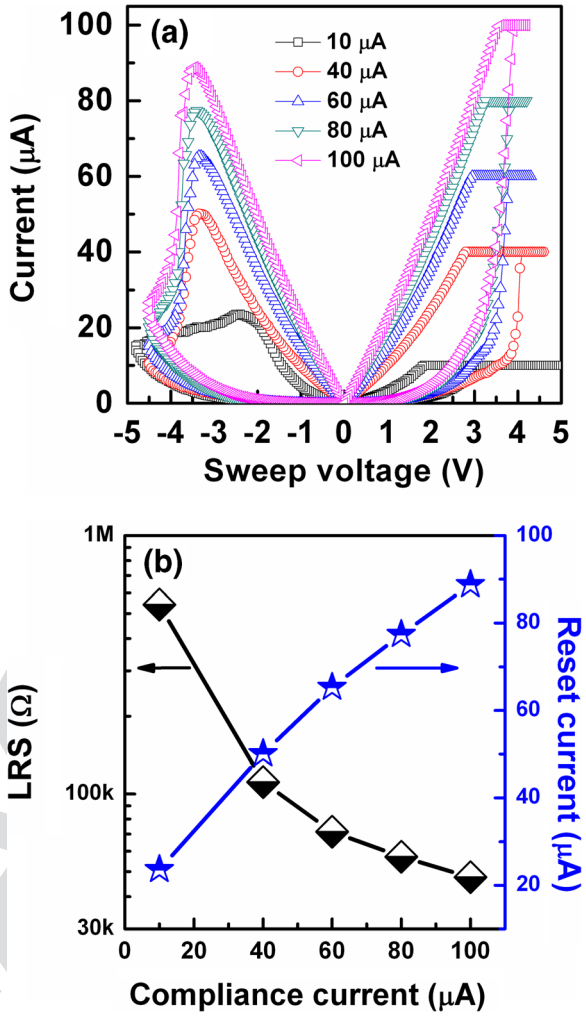

Figure 7 I-V hysteresis characteristics (a) LRS and RESET currents (b) with 10- to 100-μA CCs. A device could be operated with a low RESET current of 23 μA.

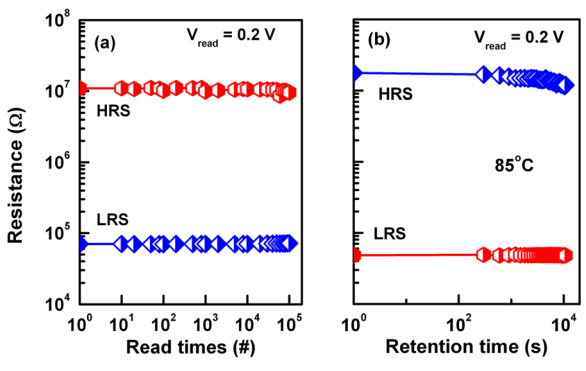

Figure 8 Long read pulse endurance and good data retention. **(a)** Long read pulse endurance of $>10^5$ cycles and **(b)** good data retention of $>10^4$ s with a good resistance ratio of $>10^2$ at 85°C are obtained at a low CC of 80 μA.

Conclusions

Improvement in resistive switching performance, particularly 10,000 consecutive switching cycles with tight distribution in LRS/HRS of $>10^2$, long read pulse endurance of $>10^5$, and good data retention of 10^4 s at 85°C, has been achieved under a low CC of 80 μA by exploiting the oxygen-getter nature of a Ti nanolayer in a W/TiO_x/TaO_x/W structure. A small device of 150 × 150 nm^2 and a defective TaO_x film are confirmed by TEM. O^{2-} ion migration rather than oxygen vacancy migration because of lower barrier height for electrons leads to a switching mechanism based on filament formation/rupture. The presence of controllable oxygen-deficient TaO_x nanofilament after switching has been investigated by AES. Furthermore, the device could be operated with a small RESET current of 23 μA. A small nanofilament diameter of 3 nm under a low CC of 80 μA has been calculated using a new method, which has a high memory density of ≈100 Tbit/$in.^2$, expected to be very useful for future sub-10-nm applications.

Competing interests

The authors declare that they have no competing interests.

Authors' contributions

AP carried out this research work under the instruction of SM. Fabrication process was also instructed by HCC and CSL. AES spectra were taken by TCT under the instruction of SM. All authors read and approved the final manuscript.

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