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Reliable resistive switching memory based on oxygen-vacancy-controlled bilayer structures†

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We fabricated resistive random access memory (RRAM) devices composed of a bilayer of AlO_x . The AlO_x layer was synthesized by atomic layer deposition (ALD) with different oxidizer sources. To control the number of oxygen vacancies, we used water and ozone as the oxidizer sources in ALD. The AlO_x layer synthesized using water as the oxidizer source ($\text{AlO}_x(\text{water})$) contained more oxygen vacancies than the AlO_x layer deposited using ozone as the oxidizer ($\text{AlO}_x(\text{ozone})$). We fabricated memory devices with a structure of $\text{Al}/\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ by ALD. After the initial forming process the devices showed reproducible and reliable RRAM characteristics. Bipolar resistive change was observed in $\text{Al}/\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ RRAM devices with low operation voltages of less than 1 V. Ohmic conduction behavior was dominant in the low resistance state and trap-controlled space-charge-limited conduction was observed in the high resistance state. The resistive switching is related to migration of oxygen vacancies and the formation/rupture of conductive filaments. Oxygen vacancies formed conductive filaments when an electric field was applied, and Joule heating broke them. The reliability of the devices was confirmed by data retention and endurance tests. The reliable bipolar resistive switching properties are attributed to a stable conductive filament in the $\text{AlO}_x(\text{water})$ layer and partial connection/disruption of a conductive filament localized at the $\text{AlO}_x(\text{ozone})$ layer.

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Introduction

Resistive random access memory (RRAM) is a promising candidate for next-generation non-volatile memory (NVM), which exploits the resistance change between high resistance state (HRS) and low resistance state (LRS). RRAM has advantages such as excellent cycling endurance, long-term retention, fast switching, low power consumption, and multistate behavior.^{1–9} Resistive switching behavior has been reported in various materials such as transition metal oxides, perovskites, metal halides, and chalcogenides.¹⁰ Recently, various oxide materials, such as AlO_x , TiO_x , HfO_x , TaO_x , ZnO , NiO_x , and CuO_x have been evaluated for use in resistive switching memory.^{11–16} Of these materials, AlO_x has great potential in RRAM applications. In AlO_x -based RRAM, conductive filaments are formed by assembly of oxygen vacancies (V_{O} s). Due to the V_{O} s, the devices exhibit the RRAM characteristic.^{17,18} To adjust the electrical characteristics of AlO_x -based RRAM, formation and assembly of V_{O} s must be controlled. Oxygen-deficient AlO_x has been achieved by chemical vapor deposition (CVD), magnetron sputtering, and pulsed laser deposition. However, those methods cannot adequately control the number of V_{O} s, or deposit them uniformly deposition on a large surface.^{19–22}

To solve these problems, atomic layer deposition (ALD) has been used to deposit stoichiometry-controlled oxide layers. ALD is based on self-limiting reaction so it has unique characteristic such as ability to deposit very uniform thin films, and to control thickness at the atomic scale. To use ALD to deposit stoichiometry-controlled oxide layers, many researchers have varied ALD conditions such as precursor exposure time, oxidizer, deposition temperature, and inert gas.^{23–25}

Moreover, to obtain reliable memory switching, a bilayer structure such as WO_x/NbO_x , AlO_x/WO_x , $\text{HfO}_x/\text{AlO}_x$, $\text{ZrO}_x/\text{HfO}_x$, $\text{TiO}_x/\text{HfO}_x$, TaO_x/WO_x , and $\text{TiO}_x/\text{TaO}_x$ has been used in the resistive switching layer in RRAM devices.^{26–34} Bilayer RRAM can provide good RRAM characteristics such as reliable and uniform electrical properties; stable resistive switching characteristics.^{26–34}

In this paper, we report use of ALD to control the amount of V_{O} s in $\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})$ bilayer structures. The first AlO_x layer was deposited using an ozone oxidizer; the second was deposited using a water oxidizer. The deposition temperature was 300 °C and the purging gas was Ar. The bottom electrode was Pt and top electrode was active Al which can induce V_{O} s in the $\text{AlO}_x(\text{water})$ layer. $\text{AlO}_x(\text{water})$ layer contains more V_{O} s than does the $\text{AlO}_x(\text{ozone})$ layer. $\text{Al}/\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ bilayer RRAM can provide reliable retention characteristic, uniform switching voltages and stable resistance switching properties compared to single oxide layer-based RRAM.

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Experimental

To fabricate the bilayer $\text{Al}/\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ structures (Fig. 1), a 10 nm Ti layer was deposited on SiO_2/Si substrate as adhesion layer by e-beam evaporation in a vacuum of $\sim 5 \times 10^{-6}$ Torr. Then a Pt bottom electrode was deposited by sputtering. AlO_x was deposited by ALD at 300 °C. The metal precursor was trimethylaluminum (TMA) and the oxidation source was water ($\text{AlO}_x(\text{water})$) and ozone ($\text{AlO}_x(\text{ozone})$). Ar was used as the purging gas during ALD. After deposition of the Pt bottom electrode 3 nm of AlO_x was deposited using an ozone as the oxidation source. Then 3 nm of AlO_x was deposited using a water as the oxidation source. Finally, Al top electrodes with diameter of 100 μm and thickness of 100 nm were deposited on the AlO_x layer by e-beam evaporation through a shadow mask. The current–voltage (I – V) characteristic, data retention, and endurance were measured using a semiconductor parameter analyzer (4200-SCS, KEITHLEY) at room temperature (RT) and atmospheric pressure, except the data retention measurement, which was also performed at 85 °C and 125 °C. During electrical measurements the Pt bottom electrode was electrically grounded, and external bias was applied to the Al top electrode.

Results and discussion

The I – V curves of the $\text{Al}/\text{AlO}_x(5 \text{ nm})/\text{Pt}$ devices exhibit a bipolar resistive switching under compliance current (CC) of 100 μA (Fig. 2). We synthesized the single layer of AlO_x by ALD under water or ozone oxidizer. The $\text{AlO}_x(\text{ozone})$ layer was more insulating than the $\text{AlO}_x(\text{water})$ layer. The $\text{AlO}_x(\text{water})$ seems to have more V_{OS} than $\text{AlO}_x(\text{ozone})$ layers; *i.e.*, that the stoichiometry of AlO_x layer can be controlled by selecting an appropriate oxidizer.^{35,36} Initial states of single layer-RRAM devices are shown in Fig. S1.† In addition, resistive switching behavior of single-layer RRAM device with $\text{AlO}_x(\text{water})$ is shown in Fig. S2.†

The I – V characteristics of $\text{Al}/\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ devices exhibit bipolar resistive switching under $\text{CC} = 100 \mu\text{A}$ (Fig. 3a). A forming process is generally needed for RRAM that exploits conductive filaments. In the forming process to form conductive filaments; a high voltage is required to overcome the high resistance state of pristine state. After the forming process, negative bias was applied from 0 to -2 V without a CC. After initial forming under positive bias, the $\text{Al}/\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ RRAM devices showed stable and uniform bipolar switching behavior. The forming voltage was observed at 1.5 V

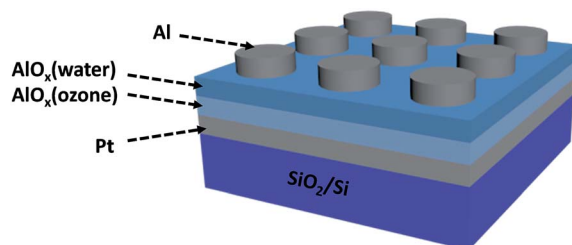


Fig. 1 Schematic of $\text{Al}/\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ bilayer structures.

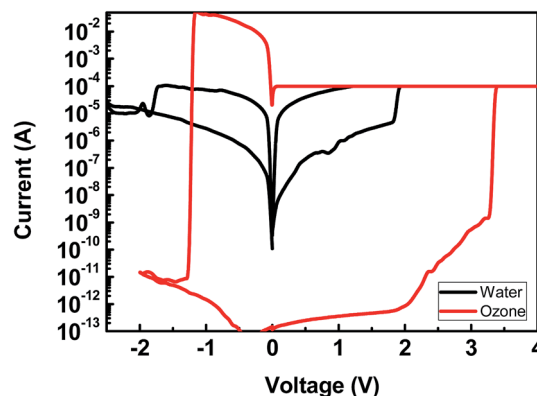


Fig. 2 I – V characteristic of $\text{Al}/\text{AlO}_x(\text{water}, 5 \text{ nm})/\text{Pt}$ and $\text{Al}/\text{AlO}_x(\text{ozone}, 5 \text{ nm})/\text{Pt}$ structure.

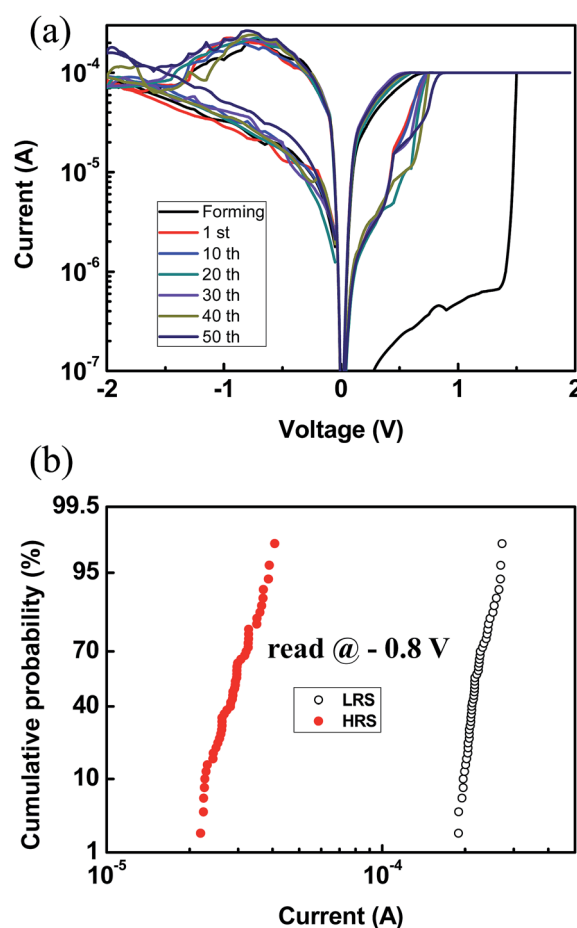


Fig. 3 (a) Typical I – V characteristics of $\text{Al}/\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ and (b) cumulative probability of current values read at -0.8 V .

and CC was 100 μA . During the second voltage sweep at positive bias, the device changed from the high-resistance state (HRS, OFF state) to a low-resistance state (LRS, conductive ON state). External bias was applied to the devices by sweeping the voltage from 0 to 2 V. When the negative bias was applied, the current declined gradually after -0.8 V and the resistance changed from LRS to HRS. After this reset process, a positive voltage was



applied from 0 to 2 V, with $CC = 100 \mu\text{A}$. Current increased rapidly at set voltage $V_{\text{set}} = 0.6 \text{ V}$. These I - V characteristics were measured under consecutive dc voltage sweeps ($0 \text{ V} \rightarrow 2 \text{ V} \rightarrow 0 \text{ V} \rightarrow -2 \text{ V} \rightarrow 0 \text{ V}$). Voltage was controlled by the Al top electrode. To investigate the distribution of HRS and LRS currents, a cumulative distribution of current was calculated using a single cell at -0.8 V (read voltage) during 50 cycles (Fig. 3b).

To determine the mechanisms of conduction and resistive switching of Al/ $\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ structured RRAM devices, a double logarithmic plot of the I - V curve was drawn in HRS and LRS (Fig. 4). At low bias ($<0.4 \text{ V}$) the slopes of the fitted lines in the double logarithmic plot were 1.001 at HRS and 1.014 at LRS; the similarity of the slope to '1' indicates that conduction behavior is ohmic ($I \propto V$). In HRS under high bias ($>0.4 \text{ V}$), the slope of this line increased to 2.792; this is the Child's law region. As the bias increased past the voltage required for the transition from the ohmic to the space-charge-limited-current (SCLC) regime, injected carriers dominated the conduction behavior.^{37,38}

To investigate the resistive switching mechanism, we measured RRAM devices with Al/ $\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ structure as above, and compared their characteristics to RRAM devices with Al/ $\text{AlO}_x(\text{ozone})/\text{AlO}_x(\text{water})/\text{Pt}$ structure (*i.e.*, AlO_x layers exchanged) while applying positive or negative bias at the top electrode. Al/ $\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ devices exhibited reliable I - V characteristic, gradual reset behavior, well-behaved bipolar filamentary switching and good uniformity under application of positive bias first (Fig. 5a), but unstable I - V characteristic, abrupt reset behavior and poor uniformity under application of negative bias first (Fig. 5b). In contrast, Al/ $\text{AlO}_x(\text{ozone})/\text{AlO}_x(\text{water})/\text{Pt}$ devices exhibited unstable I - V characteristic, abrupt reset behaviour, and poor uniformity (Fig. 5c) under application of positive bias first, and unstable I - V characteristics, gradual reset behaviour, and poor uniformity (Fig. 5d) under application of negative bias first.

To determine how the Al top electrode affects the device, we fabricated one with a Pt top electrode instead of an Al top electrode. The I - V characteristics of the Pt/ $\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ RRAM were similar to those of the Al/ $\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ devices, such as the gradual reset behavior, set

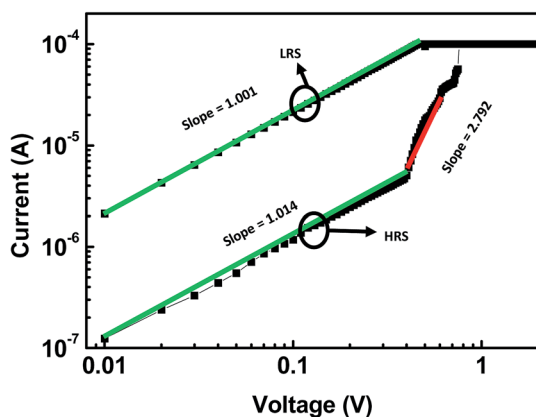


Fig. 4 Double logarithmic plot of LRS and HRS regions. Linear fitting is shown in the figure.

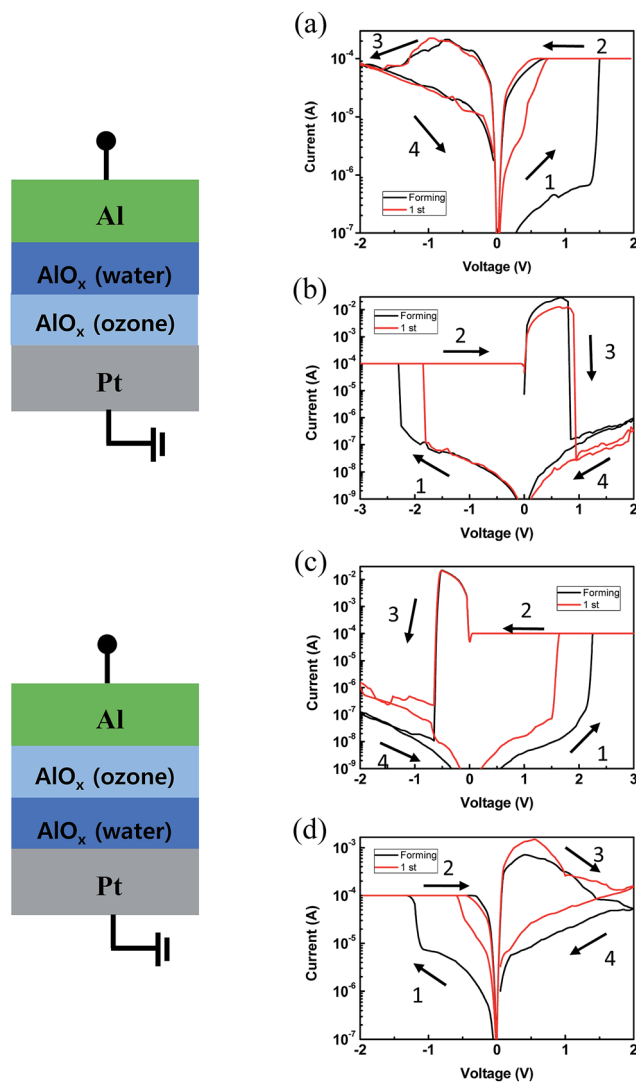


Fig. 5 I - V characteristic of Al/ $\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ with application of (a) positive bias first and (b) negative bias first. I - V characteristic of Al/ $\text{AlO}_x(\text{ozone})/\text{AlO}_x(\text{water})/\text{Pt}$ structure with application of (c) positive bias first and (d) negative bias first.

and reset voltage (Fig. 6). However, the Pt/ $\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ devices were not reliable. Only the devices with Al top electrode showed stable switching behavior.

These results suggest that the switching mechanism of Al/ $\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ devices is based on the difference in the number of V_{O} s in the two AlO_x layers. The $\text{AlO}_x(\text{water})$ layer has a large number V_{O} s, so conductive filaments remain stable, whereas the $\text{AlO}_x(\text{ozone})$ layer has few V_{O} s, the conductive filaments partially dissociate. When negative bias is applied, electrons move from the top electrode to the bottom electrode. During this process, the current density increases in interface between $\text{AlO}_x(\text{water})$ and $\text{AlO}_x(\text{ozone})$ layer, and conductive filaments partially dissociate in the top part of $\text{AlO}_x(\text{Ozone})$ layer (Fig. 7). The active Al top electrode which contacts the AlO_x layer deposited using water oxidant increases the number of V_{O} s in $\text{AlO}_x(\text{water})$. It will be very important to know the exact oxygen vacancy concentrations for understanding the exact nature of



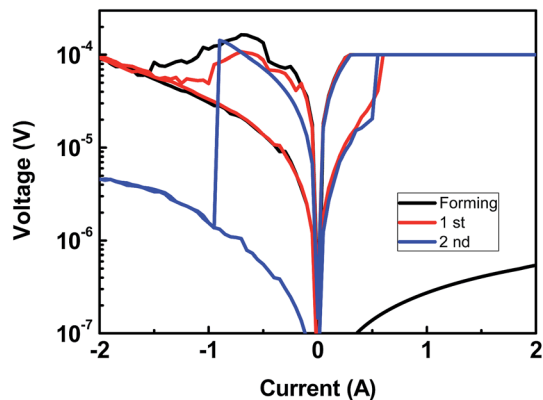


Fig. 6 I - V characteristic of Pt/AlO_x(water)/AlO_x(ozone)/Pt structure.

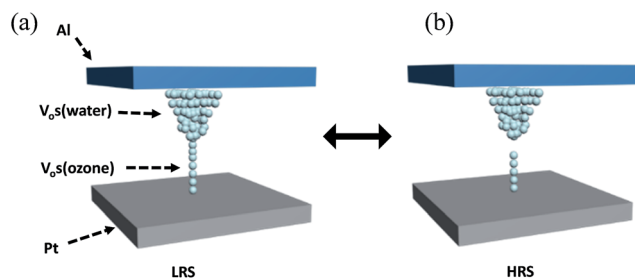


Fig. 7 Schematics illustration for switching mechanism.

defects in the resistive switching layer. We synthesized both oxide layers with the same thickness and conditions other than oxidizers (water vs. ozone), but the current-voltage response was very different. Although the exact vacancy concentration is not available it is possible to conclude that the defect concentration is very different according to oxidizers (Fig. S1†).

The cycling endurance of Al/AlO_x(water)/AlO_x(ozone)/Pt devices was measured using consecutive ac voltage pulses to verify the electrical stability under $V_{\text{set}} = +2$ V and $V_{\text{reset}} = -2$ V (Fig. 8a). The width of the voltage pulse was 10 ms and the read voltage was -0.3 V. The endurance characteristics varied slightly over time, but both LRS and HRS states retained the memory window without failure for up to 500 cycles. To estimate the non-volatile memory characteristic, data retention tests were performed at RT, 85 °C and 125 °C (Fig. 8b). Both LRS and HRS were read at -0.3 V for 10⁴ s. The current fluctuated, but the memory window was not significantly degraded at any temperature. The stable retention property at elevated temperature (85 °C and 125 °C) indicates good reliability, and shows thermal stability with constant resistance ratio. From retention and endurance characteristics, we conclude that Al/AlO_x(water)/AlO_x(ozone)/Pt RRAM devices have high uniformity and reliability.

In the Al/AlO_x(water)/AlO_x(ozone)/Pt RRAM devices, conductive filaments remain stable in the AlO_x(water) layer but the filaments partially dissociated in the AlO_x(ozone) layer because the oxide layers have different amounts of V_os. Because filaments partially dissociated in the AlO_x(ozone) layer during

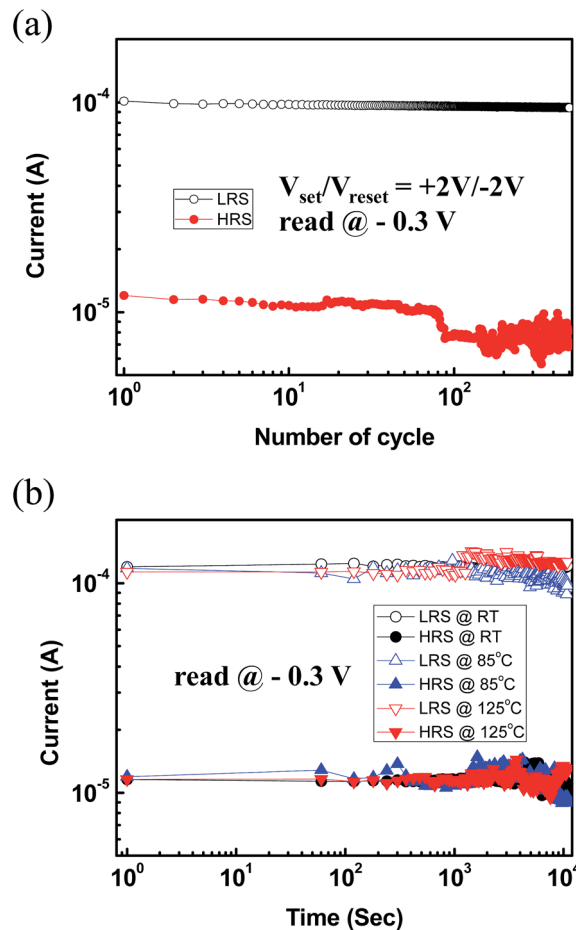


Fig. 8 Reliability of Al/AlO_x(water)/AlO_x(ozone)/Pt devices. (a) Endurance measurement. (b) Data retention properties measured at RT, 85 °C, and 125 °C.

reset process, Al/AlO_x(water)/AlO_x(ozone)/Pt RRAM devices exhibit the gradual reset behavior and immune to hard breakdown. The set and reset processes are normally affected by the competition among different filamentary paths at randomly uncertain filament sizes, orientations, and locations, so the process has a random component. However the Al/AlO_x(water)/AlO_x(ozone)/Pt RRAM devices have restricted set and reset area, resulting in good uniformity and reliability. To ensure the different amounts of V_os between AlO_x(water) and AlO_x(ozone) layer, top electrode is active Al which contacts the AlO_x(water) layer and increases the number of V_os in AlO_x(water).

In this experiment, we optimized the bilayer RRAM devices composed of Al/AlO_x(water)/AlO_x(ozone)/Pt structures; to control the number of V_os, we used two oxidizers. When water was used as the oxidizer the AlO_x layer had more V_os than when ozone was used as the oxidizer. Because the oxide layers have different amounts of V_os conductive filaments remain stable in the AlO_x(water) layer which has a large number of V_os but the filaments partially dissociated in the AlO_x(ozone) layer, which has relatively fewer V_os. Use of an active Al top electrode, can increase the number of V_os in the AlO_x(water) layer. The devices exhibit reproducible and reliable RRAM characteristic due to



partial dissociation of conductive filaments in the AlO_x (ozone) layer.

The main purpose of this work is to show the fabrication of reliable memory devices based on bilayer structure with the same oxide material by changing the oxidizer during deposition. This is very versatile and facile way to fabricate the bilayer structure with different defect concentrations. In addition, this method can simplify the fabrication processes and can effectively reduce the process cost/time.

Conclusions

We used ALD to fabricate RRAM devices composed of $\text{Al}/\text{AlO}_x(\text{water})/\text{AlO}_x(\text{ozone})/\text{Pt}$ structure. After the initial forming process, the devices exhibit reliable and reproducible bipolar resistive switching behavior. Set voltage (V_{set}) was 0.6 V and reset occurred gradually after -0.8 V. Our devices exhibit stable bipolar switching behavior, switching endurance >500 cycles, and data retention for 10^4 s at 125 °C. The conduction mechanism was ohmic in LRS and trap-controlled SCLC in HRS. The resistive switching mechanism is related to migration of V_{os} , which form conductive filaments when an electric field is applied. Joule heating breaks the filaments in $\text{AlO}_x(\text{ozone})$ layer. Conductive filaments remain stable in the $\text{AlO}_x(\text{water})$ layer which has a large number of V_{os} , whereas the filaments partially dissociate in the $\text{AlO}_x(\text{ozone})$ layer. This work can be a good solution to fabricate reliable RRAM devices since the fabrication process is simple and only single resistive switching material is needed.

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Notes and references

- R. Waser and M. Aono, *Nat. Mater.*, 2007, **6**, 833–840.
- A. Sawa, *Mater. Today*, 2008, **11**, 28–36.
- R. Waser, R. Dittmann, G. Staikov and K. Szot, *Adv. Mater.*, 2009, **21**, 2632–2663.
- H. Akinaga and H. Shima, *Proc. IEEE*, 2010, **98**, 2237–2251.
- E. Linn, R. Rosezin, C. Kugeler and R. Waser, *Nat. Mater.*, 2010, **9**, 403–406.
- M. Di Ventra and Y. V. Pershin, *Mater. Today*, 2011, **14**, 584–591.
- M. H. Lee and C. S. Hwang, *Nanoscale*, 2011, **3**, 490–502.
- 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.
- J. Park, K. P. Biju, S. Jung, W. Lee, J. Lee, S. Kim, S. Park, J. Shin and H. Hwang, *IEEE Electron Device Lett.*, 2011, **32**, 476–478.
- T. Hasegawa, K. Terabe, T. Tsuruoka and M. Aono, *Adv. Mater.*, 2012, **24**, 252–267.
- L. J. Zhang, R. Huang, M. H. Zhu, S. Q. Qin, Y. B. Kuang, D. J. Gao, C. Y. Shi and Y. Y. Wang, *IEEE Electron Device Lett.*, 2010, **31**, 966–968.
- Z. Fang, H. Y. Yu, X. Li, N. Singh, G. Q. Lo and D. L. Kwong, *IEEE Electron Device Lett.*, 2011, **32**, 566–568.
- J. W. Seo, S. J. Baik, S. J. Kang, Y. H. Hong, J. H. Yang and K. S. Lim, *Appl. Phys. Lett.*, 2011, **98**, 233505.
- Y. Wu, S. M. Yu, B. Lee and P. Wong, *J. Appl. Phys.*, 2011, **110**, 094104.
- S. Yu, H. Y. Chen, B. Gao, J. Kang and H. S. Wong, *ACS Nano*, 2013, **7**, 2320–2325.
- J.-M. Song and J.-S. Lee, *Sci. Rep.*, 2016, **6**, 18967.
- Y. Wu, S. M. Yu, B. Lee and P. Wong, *J. Appl. Phys.*, 2011, **110**.
- D. S. Jeong, R. Thomas, R. S. Katiyar, J. F. Scott, H. Kohlstedt, A. Petraru and C. S. Hwang, *Rep. Prog. Phys.*, 2012, **75**, 076502.
- S. Lee, W. G. Kim, S. W. Rhee and K. Yong, *J. Electrochem. Soc.*, 2008, **155**, H92–H96.
- S. Takata, R. Tanaka, A. Hachiya and Y. Matsumoto, *J. Appl. Phys.*, 2011, **110**, 103513.
- S. S. Lee, E. S. Lee, S. H. Kim, B. K. Lee, S. J. Jeong, J. H. Hwang, C. G. Kim, T. M. Chung and K. S. An, *Bull. Korean Chem. Soc.*, 2012, **33**, 2207–2212.
- W. Kim, S. I. Park, Z. P. Zhang and S. Wong, *IEEE Trans. Electron Devices*, 2014, **61**, 2158–2163.
- L. Assaud, M. Hanbuecken and L. Santinacci, in *Atomic Layer Deposition Applications 8*, ed. J. W. Elam, A. Londergan, O. VanDerStraten, F. Roozeboom, S. DeGendt, S. F. Bent and A. Delabie, 2012, pp. 151–157.
- J. Hu and H. S. P. Wong, *J. Appl. Phys.*, 2012, **111**, 044105.
- L. G. Wang, X. Qian, Y. Q. Cao, Z. Y. Cao, G. Y. Fang, A. D. Li and D. Wu, *Nanoscale Res. Lett.*, 2015, **10**, 135.
- S. M. Sadaf, X. Liu, M. Son, S. Park, S. H. Choudhury, E. Cha, M. Siddik, J. Shin and H. Hwang, *Phys. Status Solidi A*, 2012, **209**, 1179–1183.
- A. Prakash, S. Maikap, C. S. Lai, T. C. Tien, W. S. Chen, H. Y. Lee, F. T. Chen, M. J. Kao and M. J. Tsai, *Solid-State Electron.*, 2012, **77**, 35–40.
- R. C. Fang, Q. Q. Sun, P. Zhou, W. Yang, P. F. Wang and D. W. Zhang, *Nanoscale Res. Lett.*, 2013, **8**, 92.
- Y. Bai, H. Q. Wu, Y. Zhang, M. H. Wu, J. Y. Zhang, N. Deng, H. Qian and Z. P. Yu, *Appl. Phys. Lett.*, 2013, **102**, 173503.
- Y. Zhang, H. Wu, Y. Bai, A. Chen, Z. Yu, J. Zhang and H. Qian, *Appl. Phys. Lett.*, 2013, **102**, 233502.
- H. Zhang, N. Aslam, M. Reiners, R. Waser and S. Hoffmann-Eifert, *Chem. Vap. Deposition*, 2014, **20**, 282–290.
- C. W. Hsu, Y. F. Wang, C. C. Wan, I. T. Wang, C. T. Chou, W. L. Lai, Y. J. Lee and T. H. Hou, *Nanotechnology*, 2014, **25**, 165202.
- C. Y. Huang, C. Y. Huang, T. L. Tsai, C. A. Lin and T. Y. Tseng, *Appl. Phys. Lett.*, 2014, **104**, 062901.



- 34 C.-T. Chou, B. Hudec, C.-W. Hsu, W.-L. Lai, C.-C. Chang and T.-H. Hou, *Microelectron. Reliab.*, 2015, **55**, 2220–2223.
- 35 H. J. Lee, M. H. Park, Y. S. Min, G. Clavel, N. Pinna and C. S. Hwang, *J. Phys. Chem. C*, 2010, **114**, 12736–12741.
- 36 O. B. F. Campabadal, M. Zabala, M. C. Acero and J. M. Rafí, *Proceedings of the 8th Spanish Conference on Electron Devices*, 2011.
- 37 L. E. Yu, S. Kim, M. K. Ryu, S. Y. Choi and Y. K. Choi, *IEEE Electron Device Lett.*, 2008, **29**, 331–333.
- 38 E. Lim and R. Ismail, *Electronics*, 2015, **4**, 586–613.

