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ABSTRACT

This work addresses the thermal stability of bipolar resistive switching in yttrium oxide-based resistive random access memory revealed through the temperature dependence of the DC switching behavior. The operation voltages, current levels, and charge transport mechanisms are investigated at 25 °C, 85 °C, and 125 °C, and show overall good temperature immunity. The set and reset voltages, as well as the device resistance in both the high and low resistive states, are found to scale inversely with increasing temperatures. The Schottky-barrier height was observed to increase from approximately 1.02 eV at 25 °C to approximately 1.35 eV at 125 °C, an uncommon behavior explained by interface phenomena.

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Resistive Random Access Memory (RRAM) devices have been gaining increasing interest in the past decades as an alternative for the existing technologies in non-volatile memory applications1 and as low-device-area synaptic elements for neuromorphic computing.2 The attractiveness of RRAM lies in its great scaling potential,3 demonstrated high endurance,4 sub-nanosecond read/write speed,5,6 and potential low-power operation.7 In devices that work based on the valence change mechanism (VCM), the basic operation principle is the change of resistance through the controlled and reversible dielectric breakdown of the oxide layer, during which a defect-rich conducting path called conductive filament (CF) forms. Upon the application of appropriate voltage or current, the cell can be switched between a low (LRS) and a high resistive state (HRS). The very first breakdown (initial CF formation) is called the electroforming process; otherwise, the HRS-to-LRS transition is referred to as “set,” while the opposite process as “reset.” The biggest limitations that have to be overcome for RRAM lie in device reliability and variability. Thus, it is important to investigate and understand the stability of the performance parameters under external stress, e.g., radiation effects7 or at elevated temperatures, and to gain insight into the physical mechanisms that govern charge transport processes in the HRS as well as in the LRS. Many possible mechanisms have been proposed for conduction in both resistive states, partly due to the large variety of materials used as electrodes and functional layers in RRAM, highlighting the importance of appropriate material choice. Yttrium oxide-based RRAM is a fairly new and promising non-volatile memory8,9 and an artificial synapse candidate.10,11 Yttria was considered as an alternative to SiO2 as a gate oxide material in CMOS applications because of its high dielectric constant, large bandgap, and good thermal stability.12,13 Unlike other transition metal oxides exploited as the functional layer in RRAM, such as hafnium or tantalum oxide, for yttria the only stable stoichiometry is Y2O3.16 It has been shown that oxygen engineering of the functional layer in hafnia-based RRAM is beneficial in lowering the operation voltages due to a high oxygen vacancy density.17,18 In the case of
yttrium oxide, however, the high oxygen defect concentration is intrinsic to the crystal structure even in the stoichiometric compound, resulting in uniform resistive switching and electronic noise characteristics.\textsuperscript{11} The cubic C-type fluorite, also known as the sesquioxide or bixbyite structure of Y\textsubscript{2}O\textsubscript{3}, is inherently rich in defects as the occupation of the anion sublattice sites is only 75\%, giving rise to structural oxygen vacancies\textsuperscript{12} and making yttria an interesting material for oxygen and defect migration-based RRAM applications.

In this work, we report on the enhanced thermal stability and strongly reduced device-to-device and cycle-to-cycle variability of yttria-based RRAM. It is shown that, compared to the highly investigated hafnium oxide or tantalum oxide-based VCM-type systems, RRAM based on yttrium oxide shows very uniform switching characteristics. The operation voltages as well as the HRS and LRS current levels exhibit high uniformity, also at elevated temperatures. The observed characteristics are ascribed to the intrinsic high oxygen defect density in yttria. Additionally, the modification of the Schottky-barrier height with increasing temperature is reported. This phenomenon is of high importance as a better understanding of device physics is inevitable for the successful integration of RRAM.

The fabrication details of the devices under test (DUTs) consisting of TiN/Y\textsubscript{2}O\textsubscript{3}/Pt are described in our previous work.\textsuperscript{12} A Keithley 4200 semiconductor characterization system (SCS) was used to carry out the electrical measurements. A voltage bias was applied to the Pt top electrode while keeping the TiN bottom electrode grounded. The internal current compliance (CC) of the SCS was used to inhibit the hard dielectric breakdown of the oxide layer. A LakeShore 336 temperature controller was used for the temperature-dependent characterization. Scanning transmission electron microscopy images were obtained by using a JEOL JEM ARM-200F.

Figures 1(a) and 1(b) show the high-angle annular dark-field scanning transmission electron microscopy (HAADF-STEM) overview of the whole thin film stack and of the yttria layer, respectively. Through the high-resolution contrast, individual grains can be visualized in the polycrystalline structure of the Y\textsubscript{2}O\textsubscript{3} film. The TiN/Y\textsubscript{2}O\textsubscript{3} interface appears to be blurred, which might be due to the formation of an interfacial layer consisting of TiO\textsubscript{x} or more likely of TiO\textsubscript{x}N\textsubscript{1-x}.\textsuperscript{1,2} It is known that ohmic electrodes with high oxygen affinity are prone to oxidize upon contact with an oxide, which in turn results in the reduction of the insulating matrix.\textsuperscript{1} The formation of such an interfacial layer can assist the switching process by creating oxygen vacancies in the oxide layer and serving as an oxygen reservoir during reset. To assess the impact of temperature on the electrical behavior, six devices were pre-cycled at room temperature in ambient conditions in order to activate the resistive switching property. The measurements at elevated temperatures were carried out in vacuum. As reported in our previous work,\textsuperscript{12} the devices are electroforming-free. In Fig. 1(c), representative \textit{I} − \textit{V} characteristics are shown under DC voltage sweep operation (\textit{V\textsubscript{switch,set}} = −2.5 V and \textit{V\textsubscript{switch,reset}} = +2.5 V) at 25 °C, 85 °C, and 125 °C for one of the DUTs. The observed bipolar resistive switching behavior has been attributed to the valence change mechanism,\textsuperscript{12} where the CF formation and rupture are related to the electric field-driven motion of anions, commonly described in literature by oxygen vacancy migration. The resistive switching effect is then related to the local reduction and oxidation of the CF.

Charge transport in both resistive states has been tested for several mechanisms by the evaluation of the \textit{I} − \textit{V} curves recorded at the three measurement temperatures. These include the space-charge-limited conduction (SCLC), Poole–Frenkel emission, ionic and ohmic conduction, Schottky-emission, nearest neighbor hopping (NNH) and variable-range hopping (VRH), and well as trap-assisted tunneling (TAT).\textsuperscript{7} In previous results on yttria-based RRAM,\textsuperscript{10,12} the conduction mechanism in the LRS was identified to follow ohmic behavior, whereas the HRS was described by the Schottky-emission. This has been confirmed now also at higher temperatures as shown in Figs. 2(a) and 2(b), for the LRS and the HRS conduction, respectively. The Schottky-emission is described by\textsuperscript{7}

\begin{equation}
I = AT^2 \exp \left( \frac{\beta}{T} \right) \left( \frac{e^{g} - e}{k_B T} \right)^{1/2},
\end{equation}

\begin{equation}
\beta = \left( \frac{e^2}{4\pi e_0 \epsilon_0} \right)^{1/2},
\end{equation}

where \textit{A}, \textit{T}, \Phi_B, \textit{k}_\text{B}, \textit{e}, \epsilon_0, and \textit{\epsilon}_0 are the Richardson constant, the absolute temperature, the effective barrier height, Boltzmann’s constant, the elementary charge, the permittivity of free space, and the relative permittivity, respectively. The indicator for the formation of a Schottky-contact can be the observation of rectifying characteristics. In the investigated devices such behavior was found as under opposite bias the current-voltage curves are asymmetric. As can be seen from (1) and (2), for Schottky-emission, the \textit{ln (I/T^2)} vs \textit{V^1/2} plot has to show a linear dependence, which is fulfilled in our case for the higher voltage region [Fig. 2(b)]. Moreover, the effective Schottky-barrier height \Phi_B can also be obtained and is plotted as a function of temperature in Fig. 2(c). The effective barrier height, associated with the Y\textsubscript{2}O\textsubscript{3}/Pt interface, was found to increase with increasing temperature from \textasciitilde 1.02 eV at 25 °C to \textasciitilde 1.35 eV at 125 °C. This phenomenon has been occasionally observed in inhomogeneous Schottky-diodes\textsuperscript{24,25} and was...
usually attributed to specific interface phenomena, such as charges at the interface, surface roughness, defects, etc. These imperfections at the interface locally modify the barrier profile, which then cannot be described with a single value of the barrier height, resulting in an inhomogeneous spatial distribution of the latter. Thus, at low temperatures, the highest probability of successful electron transport is at the lowest barriers. However, with increasing temperature, the number of electrons which have sufficient energy to overcome the higher barriers increases, which in turn results in the observed increase in the effective barrier height. In the present case, a plausible reason for this inhomogeneity can be the rough Y2O3/Pt interface as visible in Figs. 1(a) and 1(b) as well as the polycrystallinity of the yttria layer showing a coexistence of two cubic phases with (222) and (400) orientation and a monoclinic (40\,/C2\,2) phase as revealed by x-ray diffraction measurements. Also, the possibility of point defects at the interface modifying the barrier profile cannot be excluded.

In Fig. 3(a), the resistance distributions of the HRS and LRS are shown as a function of temperature for all measured devices \( V_{\text{read}} = -0.2 \, \text{V} \). Note that for each device and at each temperature, at least twenty DC cycles were carried out, i.e., at least 120 measurement points are included in the statistical analysis shown in Figs. 3(a) and 3(b) for each temperature value. For both resistance states, a semiconducting behavior was observed, i.e., the resistance decreases with increasing temperature. This excludes the possibility of information storage via a metallic filament as in Conducting Bridge RAM/ Electrochemical Metallization Memories. The standard deviation of the resistance values is reduced at higher temperatures. Both resistive states exhibit good thermal stability as no enhanced degradation can be observed of either the HRS or the LRS resistance. In our previous study, we have also shown that the devices exhibit good retention characteristics in the ten year extrapolation at 85 °C and ambient pressure. In Fig. 3(b), the set and reset voltage distributions are shown cumulatively for all DUTs. The reset voltage is defined here as the voltage where the LRS current starts to decrease. We find very low operation voltages at all temperatures with \( V_{\text{set}} < 2 \, \text{V} \) and \( V_{\text{reset}} \approx 1 \, \text{V} \). Both \( V_{\text{set}} \) and \( V_{\text{reset}} \) exhibit low device to device variation as indicated by the small data spread. A higher uniformity and a shift to lower voltages with increasing temperature were observed, which is more pronounced in the case of \( V_{\text{set}} \). Similar behavior was found in TaO\(_x\) as well as in HfO\(_2\)-based devices and has been related to the increased mobility of oxygen ions and the higher probability of creating oxygen vacancies at increased temperatures. In the case of the yttria-based devices, we find that the change of the HRS and LRS resistance as well as of \( V_{\text{set}} \) and \( V_{\text{reset}} \) is strongly reduced in comparison to the hafnia and tantalum oxide-based devices. This, including the low operation voltages, can be understood to be consequential to the inherent
abundance of oxygen vacancies in yttrium oxide. As already mentioned, these so-called constitutional oxygen vacancies are intrinsic to the crystal structure and form non-intersecting channels in the oxide, which can serve as low energy ion migration paths, thus facilitating filament formation. This in turn leads to uniform switching characteristics and reduced inter- and intradevice variability.

In conclusion, the operation voltages and the resistance window in Y$_2$O$_3$-based RRAM show high thermal stability at elevated temperatures. Investigating the temperature dependence of the electrical characteristics revealed that the distribution of the set and reset voltages as well as the high and low resistive state resistances becomes narrower with increasing temperature. For all devices and test temperatures, the operation voltage window was found to span from $-2V$ to $1V$. It has been confirmed that resistive switching in the investigated devices is based on the valence change mechanism as both resistive states exhibit semiconducting behavior. The LRS charge transport shows ohmic conduction, whereas the HRS conduction is characterized by Schottky-emission. The obtained effective Schottky-barrier height was found to increase from $\sim1.02\,\text{eV}$ at $25\,\text{°C}$ to $\sim1.35\,\text{eV}$ at $125\,\text{°C}$ and has been associated with barrier height inhomogeneity at the interface.

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DATA AVAILABILITY

The data that support the findings of this study are available from the corresponding author upon reasonable request.

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