



Cite this: *Nanoscale Horiz.*, 2024, 9, 828

Received 12th December 2023,
Accepted 20th February 2024

DOI: 10.1039/d3nh00554b

rsc.li/nanoscale-horizons

Unraveling the origins of the coexisting localized-interfacial mechanism in oxide-based memristors in CMOS-integrated synaptic device implementations†

Eng Kang Koh,^{id ab} Putu Andhita Dananjaya,^{id a} Han Yin Poh,^{id ab} Lingli Liu,^{id a}
 Calvin Xiu Xian Lee,^{id ab} Jia Rui Thong,^{ab} Young Seon You^b and
 Wen Siang Lew^{id *a}

The forefront of neuromorphic research strives to develop devices with specific properties, *i.e.*, linear and symmetrical conductance changes under external stimuli. This is paramount for neural network accuracy when emulating a biological synapse. A parallel exploration of resistive memory as a replacement for conventional computing memory ensues. In search of a holistic solution, the proposed memristive device in this work is uniquely poised to address this elusive gap as a unified memory solution. Opposite biasing operations are leveraged to achieve stable abrupt and gradual switching characteristics within a single device, addressing the demands for lower latency and energy consumption for binary switching applications, and graduality for neuromorphic computing applications. We evaluated the underlying principles of both switching modes, attributing the anomalous gradual switching to the modulation of oxygen-deficient layers formed between the active electrode and oxide switching layer. The memristive cell (1R) was integrated with 40 nm transistor technology (1T) to form a 1T–1R memory cell, demonstrating a switching speed of 50 ns with a pulse amplitude of ± 2.5 V in its forward-biased mode. Applying pulse trains of 20 ns to 490 ns in the reverse-biased mode exhibited synaptic weight properties, obtaining a nonlinearity (NL) factor of < 0.5 for both potentiation and depression. The devices in both modes also demonstrated an endurance of $> 10^6$ cycles, and their conductance states were also stable under temperature stress at 85 °C for 10^4 s. With the duality of the two switching modes, our device can be used for both memory and synaptic weight-storing applications.

New concepts

This manuscript introduces a scalable CMOS-integrated memristive device designed to demonstrate dual-mode functionality encompassing both localized (digital) and interfacial (analog) switching mechanisms. These mechanisms exhibit different charge transport characteristics that cater to diverse application requirements. Previous research efforts have either primarily focused on interfacial effects as noise disruption to the filamentary switching or lacked demonstration of a scalable device viable for real-world applications. This limitation arises from the inherent challenge of effectively managing and controlling both distinct switching phenomena. The hurdle was circumvented in this work by integrating the memristive device with an NMOS transistor and leveraging the gate voltage to prevent undesired changes in the switching mechanism. Furthermore, the investigation of the memristive device switching site then revealed the bottom electrode as the active electrode, with subsequent electrode engineering enabling the facilitation of both localized and interfacial switching. The integration continued to aid in terms of enabling fast digital switching in the localized switching mode due to abrupt switching characteristics. Simultaneously, its analog switching characteristics allowed for the multiple states to be programmed, suitable for synaptic weight applications. Overall, this integration not only addresses the distinct challenges associated with dual-mode switching but also promises a paradigm shift in versatile device applications.

Introduction

The limitations of the conventional von Neumann architecture have brought about a paradigm shift in computing memory architectures. Oxide-based memristors have emerged as a fore-runner for next-generation memory due to their promising characteristics, such as high switching speed, scalability, low energy consumption, and good retention and endurance capabilities.¹ The switching properties of these memristors can be broadly classified into abrupt and gradual switching characteristics. The abrupt switching is generally reported to be driven by localized conductive filament/s, whereas the gradual switching is caused by either the formation of multiple localized filaments or interfacial effects.² In recent years, several

^a School of Physical and Mathematical Sciences, Nanyang Technological University, 637371, Singapore. E-mail: WenSiang@ntu.edu.sg

^b GLOBALFOUNDRIES Singapore Pte Ltd, 60 Woodlands Industrial Park D Street 2, Singapore 738406, Singapore

† Electronic supplementary information (ESI) available. See DOI: <https://doi.org/10.1039/d3nh00554b>



works have reported on the discovery of the gradual switching characteristics within typical memristive devices, which generally only possess abrupt switching characteristics.^{3–5} Such devices typically require specific processes or conditions to achieve each switching characteristic. Biju *et al.* demonstrated filamentary switching characteristics with a thinner switching layer (<15 nm) and gradual switching characteristics with a thicker switching layer from a WO_x-based memristor.⁶ Kim *et al.* showed the impact of annealing to obtain different switching characteristics by altering the O/Ta content in a tantalum oxide-based memristive device.⁷ Others utilized large devices of more than 1 μm², hence not showing the practical scalability of such devices.^{8–10} In this work, we report the coexistence of abrupt switching and anomalous interfacial switching characteristics within a single scalable Pt/Ta₂O₅/Ta₂N memristive device and its possible applications.

As one of the forerunners of emerging memory, the filamentary memristor aims to bridge the gap between fast but volatile memory (*e.g.*, dynamic random access memory) and non-volatile but relatively slower storage (*e.g.*, Not-AND flash). The combination of fast and data persistence eliminates the need for frequent data transfers between traditional memory and storage, enhancing processing speed and ensuring that the data remain intact in the face of system failures. This also results in increased power efficiency due to the removal of the need for frequent data refreshes to keep the data intact. The switching speed (<100 ns) and non-volatility (>10 years) of filamentary memristors make them suitable candidates for memory applications by offering advantages over both storage and memory in terms of switching speed and non-volatility.^{11,12} Concurrently, non-filamentary memristors have also been explored in neuromorphic research.^{13–15} The analog nature of such memristive devices, due to their gradual resistive switching characteristics, aligns well with the intricacies of the human brain, allowing for the creation of artificial neural networks that closely mimic the behavior of biological synapses. The synaptic weights, also known as the connection strength between the neurons, are represented by the conductance of each device, and the analog properties enable different weights to be stored *via* a step-by-step weight adjustment during the neural network training process.^{16,17} However, such devices are typically affected by their relatively poorer retention compared to their filamentary counterpart due to self-diffusion problems.¹⁸ Hence, having a single device with both abrupt and analog switching characteristics coexisting is desirable because it is promising to unite these two essential applications within a single unified memory chip. In such a memory chip implementation, our devices operating in gradual switching would be the synaptic weight, connecting the nodes from a previous layer to the next layer.^{19,20} Additionally, because the operation scheme happens after chip fabrication, the chip would be considered “field-programmable,” meaning that each core (a single crossbar array) can be assigned for various purposes, such as digital computing, storage, or neural computing. This leads to an overall improvement in cost efficiency and fabrication steps for foundries.

Results and discussion

Device structures and two operation modes

Fig. 1a shows the transmission electron spectroscopy (TEM) image of the (top to bottom) memristive device from this work. 3 nm of Ta₂O₅ and 30 nm of Pt were blanket deposited on the Ta₂N vertical interconnect access (*via*) to form the device shown in the TEM image, while the zoomed-out image was depicted in the inset. The device size can also be seen to be limited by the size of the *via*, which is about 120 nm across. It could also be observed that while the Pt and Ta₂N layers were polycrystalline, the Ta₂O₅ switching layer was amorphous. Fig S1a (ESI†) shows the energy-dispersive X-ray spectroscopy (EDS) mapping for Pt, Ta, and O to verify the deposited materials. Fig. 1b then shows a schematic of the cross-sectional of the 1T–1R device featured in this work. The area of interest highlighted in red corresponds to the inset in Fig. 1a.

Two operating modes are presented in this work, by which the devices were subjected to forward and reverse-biased modes on the same device. In this work, the ground is permanently fixed with respect to the Pt top electrode (TE). As a result, the forward-biased mode requires a positive voltage applied on the bottom electrode (BE) for the set operation and a negative voltage on the BE for the reset operation. Conversely, while in the reverse-biased mode, a negative voltage must be applied on the BE for the set operation; applying a positive voltage on the BE yielded the reset operation. Memristive devices are commonly subjected to an electrical characterization using direct current (DC) *I*–*V* double-sweep to elucidate the switching mechanism.^{21,22} A resultant *I*–*V* hysteresis loop is obtained due to the changing current response level caused by the change of potential difference across the devices. Applying the DC *I*–*V* double-sweep on the device, a forming voltage is always required to attain the forward-biased mode. The forming process is said to be needed if the current level of the subsequent cycles deviates from the first cycle. In this mode, Fig. 1c reveals the multilevel cell (MLC) switching characteristics of the memristive device. The forward-biased *IV* curve indicated by the blue arrows shows ten consecutive DC cycles of fifteen sets corresponding to a compliance current (*I*_{CC}) variation of 0.1 mA to 1.5 mA in increments of 0.1 mA. The set-sweep and reset-sweep voltages were kept constant at 1 V and –1.5 V, respectively. An abrupt jump in the current level was observed in this *I*_{CC} variation method in the forward-biased mode, in which a larger *I*_{CC} imposed led to a higher high conductive state (HCS). Imposing an *I*_{CC} prevents the device from undergoing hard breakdown while halting the movement of conductive ions within the switching layer.⁵ Additionally, the MLC could also be obtained by varying the reset-sweep voltage. Fig. 1d depicts ten consecutive DC cycles of six sets corresponding to a reset-sweep variation of –0.5 V to –1.0 V. The set-sweep voltage and *I*_{CC} were kept constant at 1 V and 1 mA, respectively. A gradual decrease in the current level was observed as the absolute value of the reset-sweep voltage increased. There was also no overlap with adjacent states for both the reset-sweep and *I*_{CC} variation methods in the forward-biased mode.



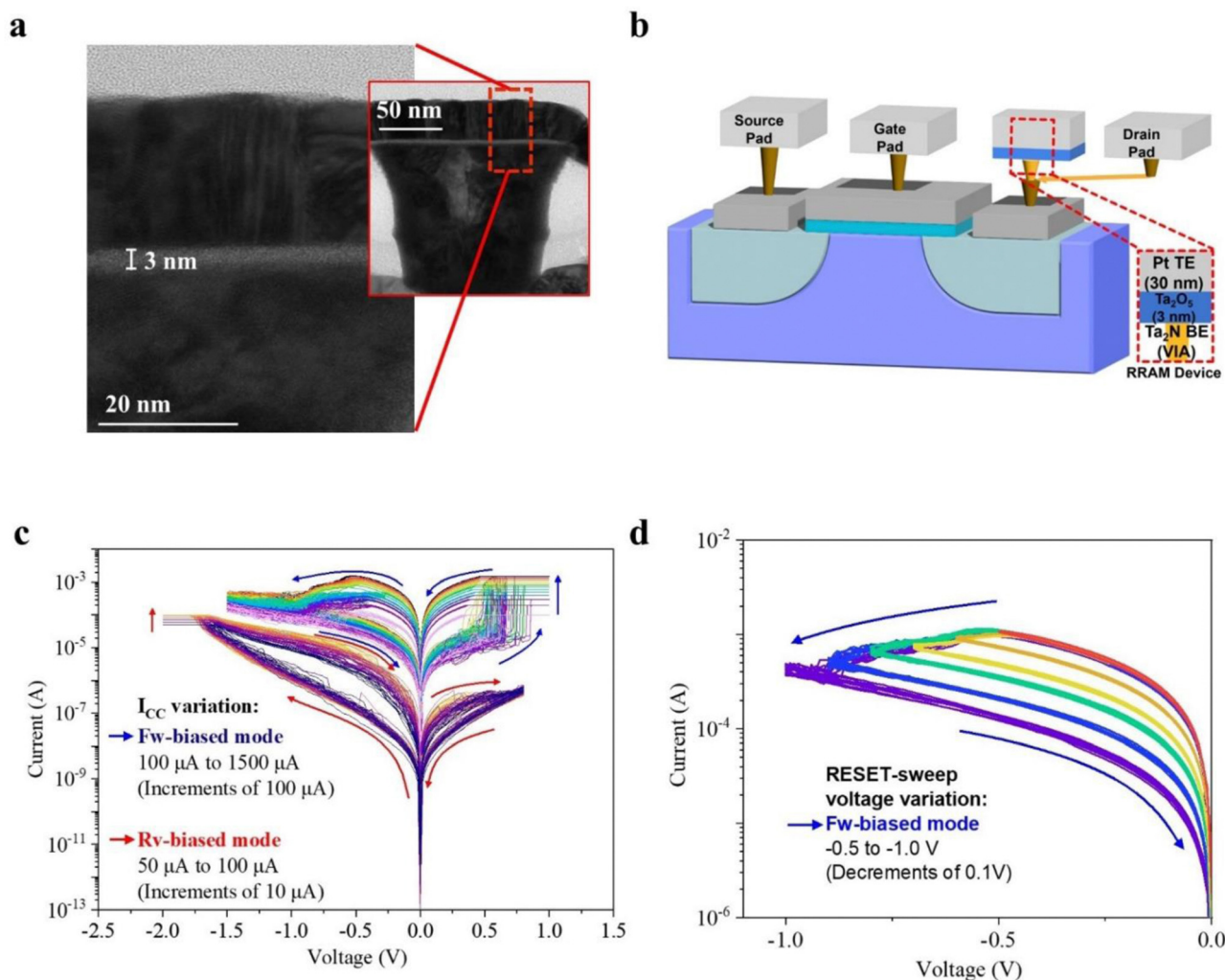


Fig. 1 (a) TEM imaging revealed the cross-sectional view of the 1R device, and the (b) schematic of the 1T-1R device (not drawn to scale). The red dotted box shows the zoomed-in cross-sectional view of the 1R scanned by TEM in (a). (c) Multilevel cell obtained by I_{CC} variation in the forward-biased (blue arrows) and reverse-biased (red arrows) mode. (d) Reset-sweep variation was also utilized to obtain a MLC in the forward-biased mode.

This indicates that the device can store the fifteen and six stable states by leveraging the external I_{CC} and reset-sweep variation, respectively. Fig. 1c also shows that the same device operated in the reverse-biased mode, as indicated by the red arrows. From a pristine state, a negative voltage was applied on the BE to set the device and, conversely, a positive voltage on the BE to reset the device. The devices are observed to not require forming to operate in its reverse-biased mode. Multilevel states were obtained by varying the I_{CC} applied on the device from 50 μA to 100 μA in increments of 10 μA , while the set-sweep and reset-sweep voltages were kept constant at -2.0 V and 0.8 V, respectively. Varying the imposed I_{CC} led to the observation of different conductance states. A higher I_{CC} applied corresponded to a higher current level (conductance levels) during the sweepback in the negative voltage region. A gradual increase in the current level in the HCS was observed as the absolute value of the set-sweep voltage increased before reaching the I_{CC} .

Elucidation of the switching mechanism in both operation modes

As mentioned previously, a forming process is required to operate the device in the forward-biased mode by applying a positive voltage (larger than the set voltage) on the BE. This forming process is also its threshold transition, enabling the device to transition from its reverse-biased mode to its forward-biased mode. This is advantageous practically (*i.e.*, when operating in pulse operation) because the switching voltage in the reverse-biased mode is unlikely to meet the forming voltage threshold during operation. Furthermore, a current range buffer exists between the two switching modes, as shown using the DC-IV curves of the devices, which are below the forward-biased mode and above the reverse-biased mode in Fig. S1b (ESI[†]). To add clarity to this, Fig. S1c (ESI[†]) shows the forming voltages of ten different devices featured in this work. Their forming voltages varied from 1.04 V to 1.52 V, with a median value of 1.32 V. The red dotted line depicts the reset voltage in



the reverse-biased mode. Since practical applications require demonstration of pulse operation, the marginal difference between the reset voltage in the reverse-biased mode and the forming voltage in the forward-biased mode becomes crucial, as a larger margin prevents unintended transition into the forward-biased mode as long as the reset voltage in the reverse-biased mode does not increase above the forming voltage threshold. This is unlike in most previous studies, in which the memristive hysteresis loop of the gradual switching sits directly above that of the abrupt switching, leading to an absence of the aforementioned current buffer.^{9,23–28} In other words, in those works, the HCS of the reverse-biased mode is also the low conductive state (LCS) of the forward-biased mode. This would lead to an undesired change in modes during pulse operations, rendering them useless in such applications. Since the devices have distinct differences in current responses, there is a need to understand the physics behind the switching behavior of the different operational modes. An experiment was designed to investigate the influence of the memristive

device size in the different modes on the switching current levels.

Fig. 2a depicts the HCS and LCS of the devices in the forward-biased mode (blue) and reverse-biased mode (red) in terms of their device size diameter. For the forward-biased mode, the LCS and HCS were obtained with a reading voltage of 0.1 V after a set sweep of 1 V ($I_{CC} = 1$ mA) and a reset sweep of -1 V, respectively. For the reverse-biased mode, the LCS and HCS were acquired by a reading voltage of 0.3 V after a set sweep of -2 V ($I_{CC} = 100$ μ A), and the HCS was obtained after a reset sweep of 1 V, respectively. The measurement was done over five cycles using five devices each to obtain the mean conductance, and the conductance ranges were then plotted as an interval plot. The results showed that the conductance states in the forward-biased mode are independent of their device sizes. This is shown by the conductance level being relatively constant regardless of the device size. This differs from the reverse-biased mode, whereby the conductance states positively correlate with the size of the devices, *i.e.*, the larger the device

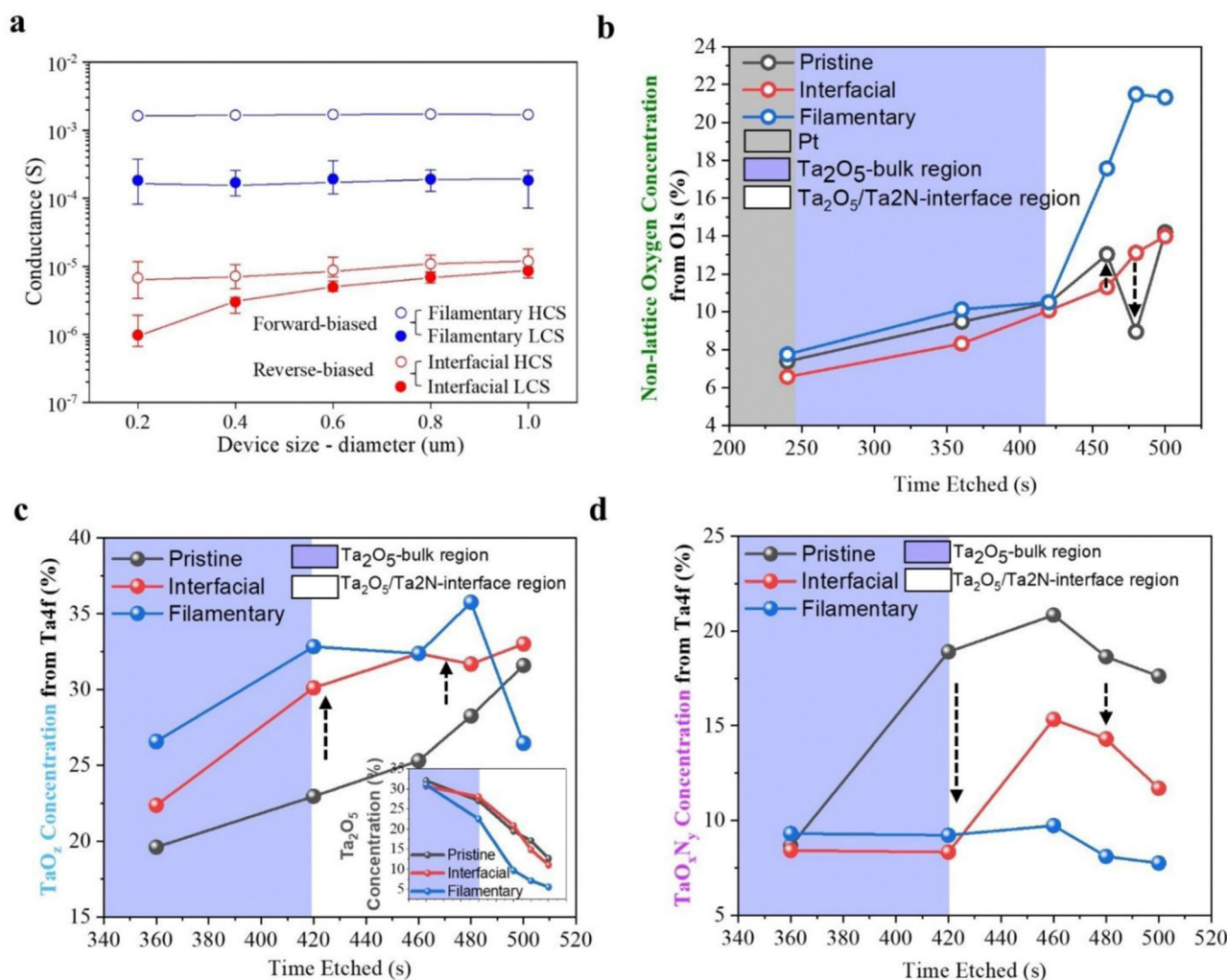


Fig. 2 (a) Conductance-size dependence study conducted on the memristive devices shows size independence in the forward-biased (blue) modes and size dependence in the reverse-biased mode. Percentage concentration of (b) non-lattice oxygen from O1s, (c) TaO_x from Ta4f, and (d) TaO_xN_y from Ta4f as obtained by XPS depth profiling (raw data and fittings presented in Fig. S2 and S3, ESI†)

area, the higher the conductance states (and also a smaller on-off ratio). Consequently, this suggests that in the forward-biased mode, resistive switching happens locally (due to filament formation). In contrast, in the reverse-biased mode, the device undergoes homogenous switching, which occurs interfacially in the reverse-biased mode. The apparent subsequent step was to identify the location of this interfacial switching through the use of X-ray photoelectron spectroscopy (XPS) depth profile analysis.

However, due to the scanning limitations of XPS, additional devices fabricated were scaled up in terms of the switching layer thickness (to 35 nm) and device diameter (100 μm). This exact size was chosen because the device switched in the same current range as the original devices in the reverse-biased mode. Three different devices (pristine – sample A [black], interfacial HCS – sample B [red], and filamentary HCS – sample C [blue]) were investigated for their switching mechanism. The fitting of the raw data is presented in Figures S2 and S3 (ESI[†]) for O1s (528–535 eV) and Ta4f (20–31 eV) peaks, respectively.^{29–33} The O1s spectra were deconvoluted into two peaks, and Ta4f was deconvoluted into five doublet peaks, respectively.^{29–33} In terms of the depth profile, there were approximately two regions: <420 s time etched for the Ta₂O₅ bulk region and the >420 s time etched for the Ta₂O₅/Ta₂N interface region. Fig. 2b shows the concentration percentage of the non-lattice as a percentage of total O1s core level peaks. It could be observed that the level of non-lattice oxygen concentrations increased at relatively the same rates from ~7% to ~10% in the bulk region for all three samples. However, this is drastically different in the interface region. Comparing samples A and B, the non-lattice oxygen concentration in sample A dipped to the bulk levels (~9%) at 480 s etch depth. Conversely, the non-lattice oxygen concentration in sample B continued to increase at the interface until the levels of both samples A and B equalized at 500 s etch depth. This suggests that a depth exists within the interface by which the percentage of lattice oxygen is higher than the depth above and below it. In other words, this is also caused by the redox reaction that happens at the Ta₂O₅/Ta₂N interface. This hypothesis is also further corroborated by TaO_xN_y concentration and TaO_xN_y concentration as a percentage of Ta4f peaks in Fig. 2c and d, respectively. In Fig. 2c, TaO_z was detected earlier and had a larger concentration in sample B compared to sample A. Similarly, in Fig. 2d, TaO_xN_y was detected earlier and had a higher concentration in sample A compared to sample B. Moving on to the analysis of sample C, observations made in Fig. 2b revealed that the non-lattice oxygen concentration percentage in sample C was similar to that of samples A and B in the bulk. However, nearer to the interface, there was a large increase in this non-lattice oxygen concentration percentage. Fig. 2c then shows that sample C contained relatively higher TaO_z (Ta sub-oxide) content percentage-wise compared to samples A and B. This also supported the revelation of the consistently lower Ta₂O₅ percentage concentration in sample C compared to samples A and B, suggesting a larger amount of oxygen vacancies or defects present throughout the switching layers of the device. Finally, it can also be seen in Fig. 2d that the TaO_xN_y concentration

percentage in sample C remained negligible compared to samples A and B.

In order to further clarify the explanation made from the XPS results in Fig. 2, Fig. 3 was created to depict the proposed switching mechanism of the memristive device in the different modes. Within the literature on the coexistence of filamentary and interfacial switching in memristive devices, the underlying mechanism behind most interfacial switching was proposed to stem from the trapping and de-trapping of charges at the oxide-inert electrode interface, for which the evidence is lacking in some prior works.^{8,23,24} In this work, we propose that for the Pt/Ta₂O₅/Ta₂N device presented, the interfacial switching happens at the oxide-active electrode interface instead. Fig. 3 illustrates that the interfacial switching of the device in the reverse-biased mode is caused by external electrical stimuli, leading to the thickness modulation of the oxide-active electrode interface. Initially, Ta₂N/Ta₂O₅/Pt layers were deposited. The redox reaction then spontaneously started to proceed at the Ta₂O₅/Ta₂N interface due to the oxygen-gettering tendencies of the tantalum nitride electrode.^{34,35} The XPS results in Fig. 2 suggest that in the reverse-biased mode, the Ta₂O₅ layer acts as an oxygen reservoir to enable the redox reaction with the Ta₂N layer. The externally applied electric field modulates the TaO_xN_y and TaO_z concentration, sandwiched by the oxidation of the Ta₂N layer below and the reduction of the Ta₂O₅ switching layer. Additionally, the Ta ions from the Ta₂N BE are also expected to contribute to the downward growth of the TaO_xN_y layer due to their upward migration from the Ta₂N electrode layer, as shown in a prior study that they are mobile in the Ta₂O₅ layer and are believed to mediate switching.³⁶ Applying a negative voltage (in the reverse-biased mode) onto the pristine device to set it resulted in the generation of non-lattice oxygen (and, in its place, oxygen vacancies), providing the device with a higher conductance. Conversely, applying a positive voltage brings these oxygen ions back down into the interface, thus increasing the rate of recombination of the oxygen vacancies and thereby reducing the device conductance. This explanation is similar to Baek *et al.*'s work on the TiN/PCMO/Pt memristor stack, where an *in situ* TEM was used to prove that the resistive switching was a result of the shrinking and expansion of the TiO_xN_y interfacial layer.³⁷ For the forward-biased mode, applying a relatively larger positive voltage causes dielectric soft-breakdown. Consequently, it introduces more defects in the form of localized oxygen vacancies throughout the switching layer (also known as the conductive filament). The filamentary switching in the forward-biased mode can be explained by the widely agreed filamentary formation and dissolution model.^{5,38–40} Previously, in Fig. 1, two typical approaches are implemented to demonstrate the multi-bit capability in the forward-biased mode, *i.e.*, current control during the set operation and voltage control during the reset operation. For the reset operation, as seen in Fig. 3, the entire filament (composed of oxygen vacancies, V_O) disconnected from the TE, resulting in a decreased current level. Subsequently, as the voltage magnitude increased negatively, the filament retracted further from the cathode, causing an additional decrease in the current level.



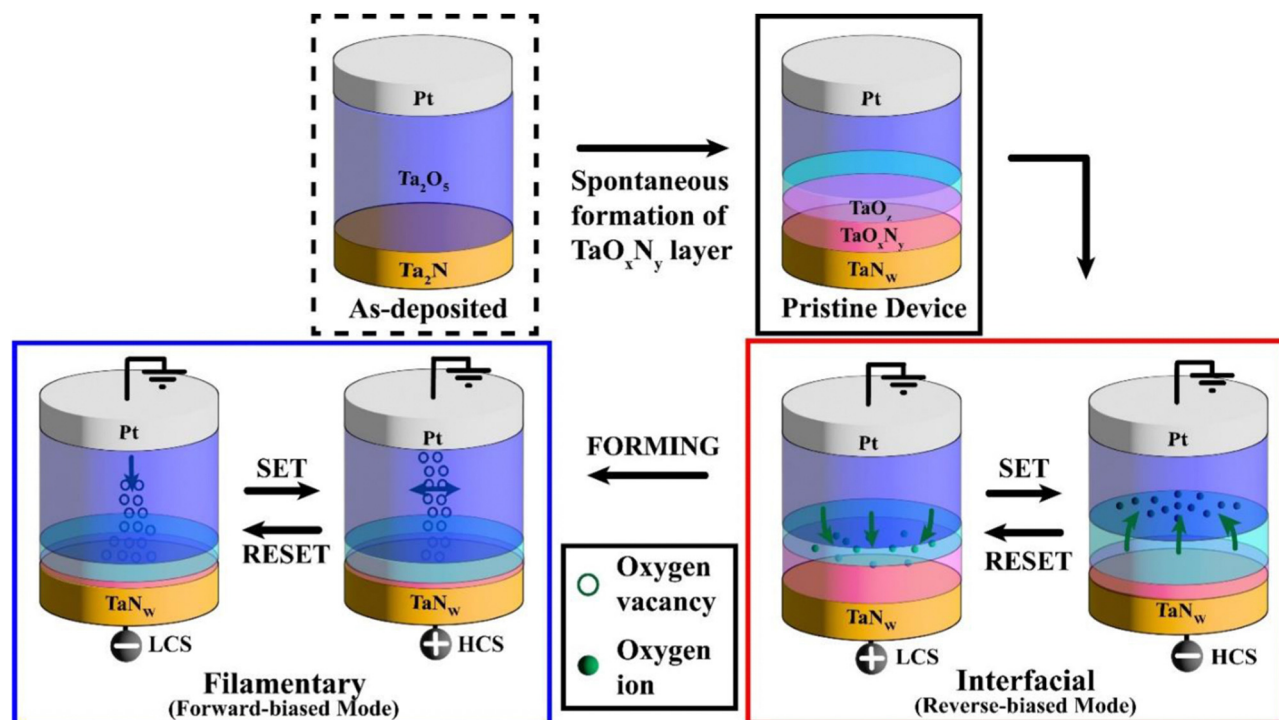


Fig. 3 Schematic diagram of the Pt/Ta₂O₅/TaN memristor switching mechanisms in both forward-biased (localized filamentary) and reverse-biased (interfacial) modes. Filamentary switching occurs when the voltage applied across the electrodes controls the migration of oxygen vacancies acting as highly conductive pathways for electrical current flow. The interfacial mode is proposed to happen due to the expansion and contraction of the spontaneously formed TaO_xN_y dominant layer within the memristive device due to voltage, effectively reducing and increasing device conductance. The transition from interfacial switching to filamentary switching requires a forming process, during which the switching layer undergoes soft-break down and oxygen vacancies are introduced.

During the set operation, a fine localized filament grows from the BE to TE and gradually expands radially as the current increases. Imposing I_{CC} then allowed for control to obtain multilevel switching, depending on the radial size of the filament, in which larger conductive filament thickness reflects a higher device conductance.

Conduction mechanism of the devices in both operation modes

Besides the switching mechanism, it is also crucial to understand the charge transport dynamics within the device during the electrical characterization (*i.e.*, the DC sweep). For example, Ohmic conduction is crucial for executing matrix-vector multiplication by harnessing Kirchhoff's Law, effectively reducing the IR drop in a large array. This effect has also been analogous to the intricacies of the functions of the biological brain.⁴¹ Conduction mechanism models reported in the literature are broadly categorized into electrode-limited or bulk-limited.⁴² The consensus with such analysis is that although there tend to be multiple mechanisms concurrently happening with the device, the fitting to various models only shows the most dominant effect.⁴² Determining the most dominant mechanism would possibly help develop new applications to exploit the memristor's switching properties during fabrication or tuning. Fig. 4 shows the conduction mechanism of the device, whereby the fitting was done on the device operated in both the forward-biased (Fig. 4a and b) and reverse-biased (Fig. 4c and d)

modes. Fig. 4a and b depict the forward-biased IV curves fitted with the Ohmic and Schottky conduction mechanisms, respectively. The insets in these figures show the region/s from which the IV data were collected. Voltage sweeps were applied across the device five times consecutively to ensure that the change in current did not occur due to the change in the conductance state of the device. The gradient and y-axis intercept were then obtained from the average fittings. Generally, the LCS low-field regions showed an ohmic conduction mechanism in both polarities in the range of [0 to 0.2 V]. This was revealed through a double arithmetic fitting with an average gradient of 1.05 and 1.07 for positive and negative polarities, respectively. Under a low field, only a small number of carriers are generated through thermal excitation. The data points in the LCS high-field region deviated from the fittings; hence, other conduction mechanism models were fitted instead. The Schottky model (shown in eqn (1)) was the best fit through a $\ln I$ against $V^{1/2}$ fitting for the data points in that region. When the device is subjected to a relatively higher electric field, an increase in thermal energy further excites the electrons, allowing them to overcome the Schottky energy barrier at the interfaces to enter the dielectric layer. This happens until a point where the set voltage is reached, and the device undergoes an abrupt resistive state switch into the HCS. Experimental dielectric constant values of 37.2 and 47.7 were obtained for the positive and negative sweeps, respectively, and the Schottky barrier height was



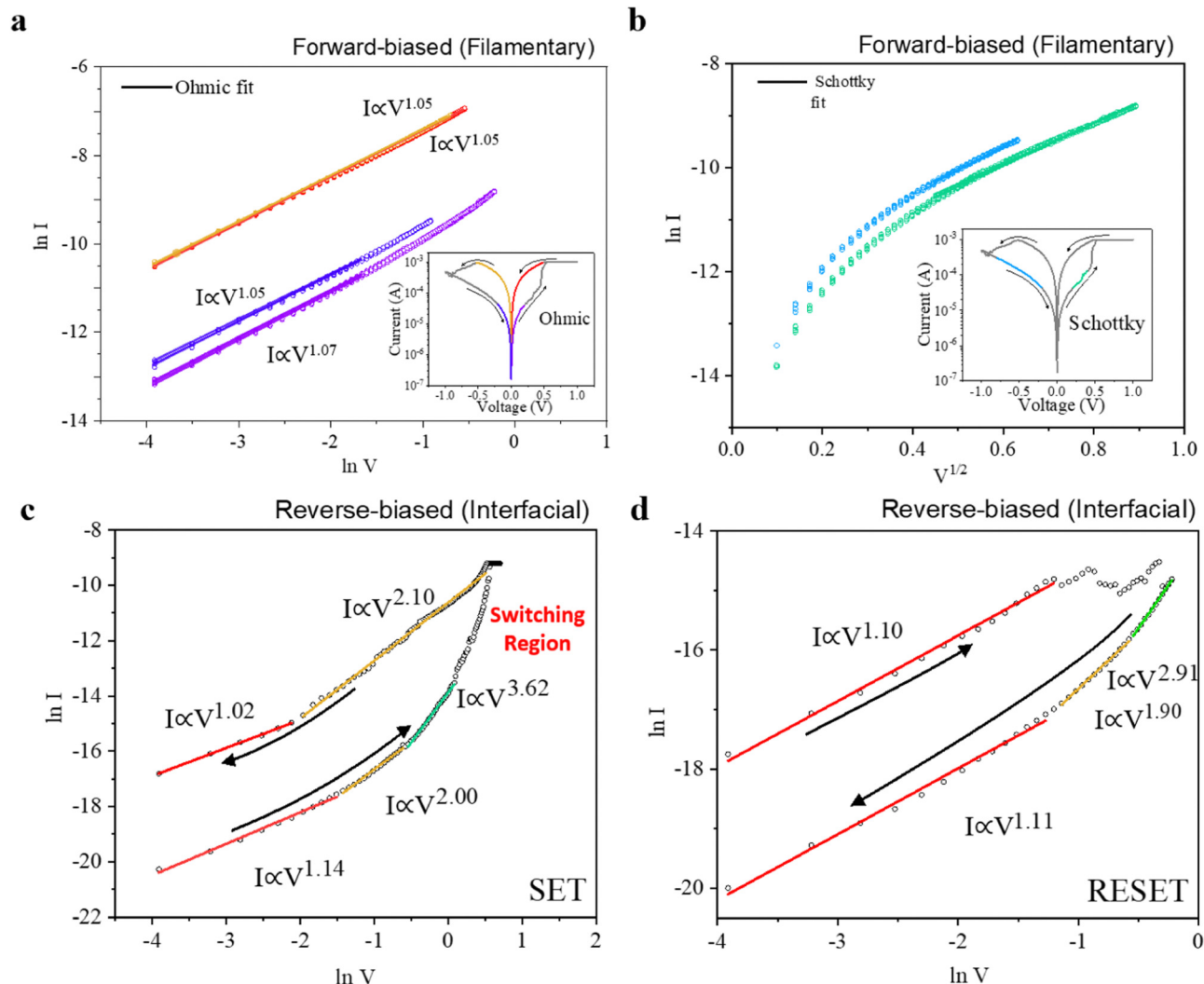


Fig. 4 Fitting and analysis of the DC I - V curve using a double logarithmic plot forward-biased mode indicated that it is dominated by (a) Ohmic mechanism in the low-field LCS and HCS, and (b) Schottky mechanism in the high-field LCS. The I - V characteristic device in its reverse-biased mode indicates an SCLC conduction mechanism for the (c) set process and the (d) reset process, in which all of the fitting ones were with adjusted R^2 values of more than 0.99.

2.09 eV for both the $\text{Ta}_2\text{N}/\text{TaO}_x\text{N}_y$ and $\text{Ta}_2\text{O}_5/\text{Pt}$ interfaces. For both positive and negative polarities in the HCS region, the double logarithmic fitting was again the best fit, with an average gradient of 1.05 in both regions. This low field coincides with the read voltage, which is beneficial when the devices are in an array-level in-memory computing implementation, and that is to avoid non-idealities like the sneak path or leakage current when dealing with positively non-linear loads (where the current through the device increases more than being proportionate with an increase in voltage).⁴³

$$J_{\text{SE}} = \frac{4\pi q m^* (kT)^2}{h^3} \exp \left[\frac{-q \left(\Phi_{\text{B}} - \sqrt{\frac{qE}{4\pi\epsilon}} \right)}{kT} \right] \quad (1)$$

Fig. 4c and d then depict the set and reset IV curves in the reverse-biased mode, respectively. In Fig. 4c, increasing the

voltage in the reverse-biased LCS region, the slope of the \ln - \ln curve evolved from a gradient of 1.14 in the low-field region (0 to -0.22 V, red) to a gradient of 2.00 (-0.22 V to -0.54 V, orange), and then finally to a gradient of 3.62 in the high-field region (> -0.54 V). When the voltage was swept back, the current response initially gave a gradient of 2.10 and subsequently back to 1.02 in the low field. The fitting in Fig. 4d then showed a similar trend in the reset operation, in which, from its LCS, the curve gradient started from 1.10, and then a decrease in current was eventually observed at -0.28 V. The sweep back at a higher field of > -0.56 in the LCS led to the gradient undergoing a transition from 2.91 from a high-field region to a gradient of 1.90 (-0.28 V to -0.56 V), then finally to a gradient of 1.11 in the low field region (0 to -0.28 V). The discussed results suggest that the space-charge limited conduction (SCLC) mechanism was the dominant conduction mechanism when operated in the reverse-biased mode, in which the slopes of the graph at different regions correspond to the Ohmic law



($\sim I \propto V$) at low field, a transition to the Mott–Gurney law ($\sim I \propto V^2$), and finally a steep increase in current injection at higher fields ($\sim I \propto V^{>2}$).⁴² To explain from the LCS, when the device was subjected to a low field in its LCS, it was in the ohmic state. In this range, the density of the thermally generated free carriers within the switching layer was larger than that of the injected carriers. As the voltage was then increased, the traps within the switching layer started to fill up, and the slope of the curve was interpreted to follow the Mott–Gurney law. These traps were fully filled as the device was swept with a high field. This correlated with an accelerated increase of the current level whereby further injected carriers were freely moving within the layer.⁴⁴ For all the best-fit lines done in this work, adjusted R^2 values were always more than 0.99.

From fitting the Schottky emission model in eqn (1) to Fig. 4b, the gradient and y-axis intercept were extracted to be ~ 4.47 and ~ -12.3 for the positive region, and ~ 3.95 and ~ -12.3 for the negative region. The dielectric constants and Schottky barrier

heights were derived from those values. Depending on the deposition methods and parameters, Ta_2O_5 thin films generally have a dielectric constant around the range of 25; however, they can go upwards to 110, as reported by Lin *et al.*^{45,46} On the other hand, TaO_x has a dielectric constant that is generally lower than Ta_2O_5 . In this work, we obtained an experimental value of ~ 37.2 and ~ 47.7 for the positive and negative sweeps, respectively. This suggests that the effective dielectric constant of the dielectric layer, which was sandwiched by the electrodes, decreased during the set operation and increased during the reset operation. Referring back to the filamentary switching mechanism schematic in Fig. 3, the percentage of TaO_x is increased due to the generation of non-lattice oxygen during the set process and decreased due to the recombination of these non-lattice oxygen species. This would lead to a drop and increase in the effective dielectric constant, respectively, consistent with TaO_x having a lower dielectric constant than the more stoichiometric Ta_2O_5 , as mentioned previously. Fig. 5a summarises the findings using

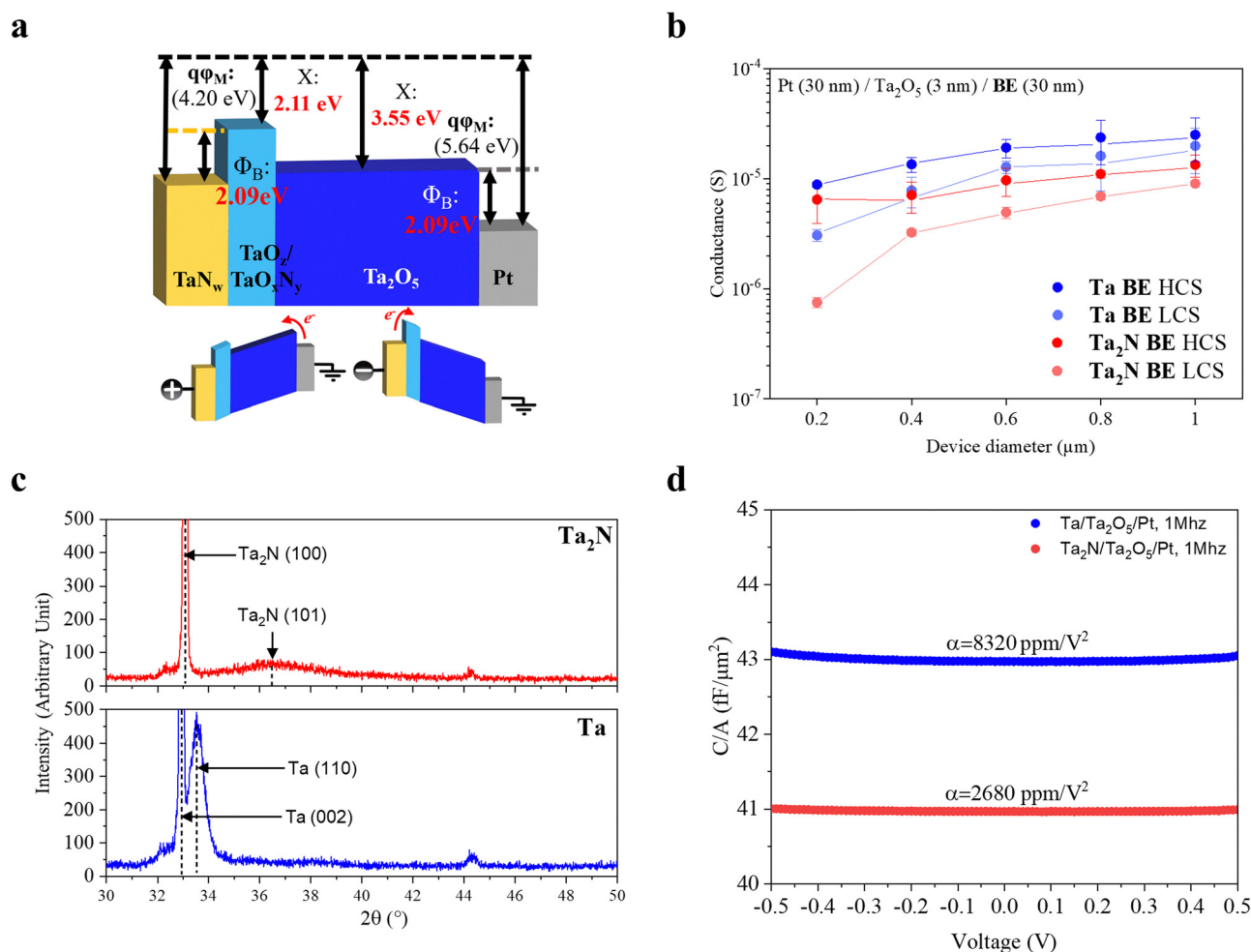


Fig. 5 (a) Schematic band diagram of the deposited Pt/ Ta_2O_5 / Ta_2N memristor. (b) Comparison of the effects of changing the BE from Ta_2N (red) to Ta (blue). Investigation of the LCS and HCS of 5 devices for each device diameter. Ta BE devices constantly had a device conductance lower than that of those with the Ta_2N active BE. (c) XRD pattern of Ta_2N and Ta deposited on a SiO_2 substrate, showing that the Ta deposited was one of a beta phase, with a dominant (002) peak and a relatively weaker peak at (110). Reactive sputtering with nitrogen yielded a strong Ta_2N peak at (100) and a weak peak at (101). (d) $C-V$ characteristics of Pt/ Ta_2O_5 / Ta_2N and Pt/ Ta_2O_5 /Ta.

an energy band diagram, in which the values in red were calculated based on this experiment, and those in brackets were both polarities, which were calculated to be 2.09 eV. Using these values, the electron affinity of the TaO₂/TaO_xN_y and Ta₂O₅ layers was determined to be 2.11 eV and 3.55 eV, respectively. Therefore, the evidence suggests the existence of Schottky barriers at both interfaces as obtained from the literature.^{47,48} As mentioned previously, eqn (1) was utilized to obtain the Schottky barrier height of the devices in both polarities, which was calculated to be 2.09 eV. Using these values, the electron affinity of the TaO₂/TaO_xN_y and Ta₂O₅ layers was determined to be 2.11 eV and 3.55 eV, respectively. Therefore, the evidence suggests the existence of Schottky barriers at both interfaces.

To provide further elucidation on the conduction and switching mechanisms mentioned above, it is necessary to begin from the pristine device shown in Fig. 3. When a negative biased (reverse-biased mode, Fig. 4c) is applied to the device, it undergoes SCLC, a bulk-limited conduction mechanism, due to which as the applied voltage across the device is increased negatively, the rate of increase of current also increases. This results in a decrease of resistance across the Ta₂O₅ bulk, with the majority of the voltage being shifted to the TaO_xN_y/TaO₂ interface, where switching is proposed to commence. This effect is shown in Fig. S4 (ESI[†]), where the *I*-*V* curve starts to exhibit a hysteresis above $> \sim 1.0$ V (as shown in Fig. S4, ESI[†]). This hysteresis gets larger as we continue to increase the said voltage. Subsequently, during the forming process to transition the device to the filamentary forward-biased mode, the oxygen vacancies form a low-resistance filament in the Ta₂O₅ bulk, connecting the two electrodes. This is where the device conduction starts to be governed by the Schottky mechanism, an electrode/interface-limited conduction mechanism.

Besides the switching layers having a direct impact on the switching characteristics, there is also much evidence of the electrode engineering affecting the device switching characteristics.^{49–53} Hence, it was also important to delve into how the nitrogen content in the BE helps with the switching characteristics of the proposed Pt/Ta₂O₅/Ta₂N devices. Fig. 5b shows a conductance plot for the memristive devices with Ta₂N (red) and Ta (blue) as the BE, operated in the DC reverse-biased mode. Five devices were measured for each device size, and the conductance variations were accounted for as error bars within each data point. The devices were switched with a set and reset voltage of -2 V and 1 V, respectively, and an I_{CC} of $100 \mu\text{A}$ was imposed during the set operation. The devices' states were read with a reference voltage of 0.1 V. It was then observed that the respective HCS and LCS are consistently higher for the Ta BE devices than for the Ta₂N BE. This could be attributed to either the nitrogen content in the BE contributing to the series resistance of the memristive device, or the difference in oxygen affinities of the different BE layers altering the oxygen content of the Ta₂O₅ switching layer directly above, or both. X-ray diffraction (XRD) was subsequently performed on 30 nm of both Ta and Ta₂N, and its results are depicted in Fig. 5c. The BE materials were deposited *via* sputtering onto SiO₂ substrates to confirm the BE material as claimed. It was then observed in the

resultant XRD patterns that the Ta that was deposited by sputtering had a strong β -Ta 002 peak at around 32.9° and a relatively weaker α -Ta 110 peak at around 33.5° .⁵⁴ Comparatively, the sputter-deposited Ta₂N had a strong hex-Ta₂N 100 peak at 33.0° and another relatively weak hex-Ta₂N 101 peak at 36.8° .⁵⁵ After observing that the film deposited by reactive sputtering was that of a hex-Ta₂N, capacitance-voltage (*C*-*V*) measurement was conducted on the Pt/Ta₂O₅/Ta and Pt/Ta₂O₅/Ta₂N memristive devices, for which the results are presented in Fig. 5d. The *C*-*V* analysis was conducted by fitting the second-order polynomial *C*-*V* equation to obtain the voltage coefficient of capacitance (VCC) shown in eqn (2):^{56–58}

$$\frac{C(V) - C_0}{C_0} = \alpha V^2 + \beta V \quad (2)$$

here, C_0 refers to the capacitance zero bias voltage, and α and β are the quadratic and linear VCC, respectively. The quadratic α term in eqn (2) is desired for analysis here, as it is known to correlate with the spontaneous formation of the interfacial layer between the BE and the switching layer. Correspondingly, this also establishes a positive relationship between the quadratic α term and the oxygen affinity of the BE. The *C*-*V* measurements in this work were all performed at 1 MHz in the range of -0.5 V to 0.5 V because it was observed that any increase in the range would cause a breakdown of the device. The C_0 of the Pt/Ta₂O₅/Ta (43.0 fF μm^{-2}) device was higher than that of the Pt/Ta₂O₅/Ta₂N devices (41.0 fF μm^{-2}). The *C*-*V* curve was fitted to eqn (2), and this yielded α values of 8320 ppm V^{-2} and 2680 ppm V^{-2} for the memristive devices with Ta and Ta₂N BE, respectively. This shows that the oxygen affinity of the Ta BE is higher than that of Ta₂N BE, which in turn suggests that the TaO_xN_y content should be higher in Ta BE devices due to the higher occurrence of oxidation in those devices compared to the Ta₂N BE memristive devices. Subsequently, resistive switching characterization was conducted on the devices with different BEs. It was then observed that the memristive devices comprising the Ta₂N BE could not be reset after the forming process; in other words, they could only be switched interracially. This would seem to conflict with literature reports of filamentary resistive switching Pt/Ta₂O₅/Ta memristive devices. However, those resistive switching layers in such reports are relatively thicker (> 3 nm), and device sizes were larger (> 100 nm diameter) compared to those used in this work.^{59,60}

1T-1R integration of the Ta₂O₅-based memristor and performance

The Ta₂O₅ memristive device in this work was then integrated into the N-channel metal-oxide semiconductor field effect transistor (NMOS) transistor at its drain. Integrating the 1T-1R cell structure allows data to be stored using the memristor element while simultaneously using the NMOS transistor to control the current across the memristive device, thus reducing the sneak current when presented in an array. In this work, the NMOS transistor is also exploited to demonstrate the control of the multilevel cell properties of the memristive devices during



practical (pulse) operations. Referring back to the 1T-1R schematic in Fig. 1b, in the forward-biased mode of the 1T-1R cell, the set process was conducted by applying a positive voltage on the source pad, and the reset operation was conducted by applying a positive voltage on the TE pad. Conversely, the reverse-biased mode required a positive voltage on the TE pad for the set operation and a positive voltage on the source pad for the reset operation. Consequently, the reference voltage and current will be with respect to the source of the transistor for the presentation of this work below (and TE considered grounded). Positive voltage was also applied on the gate pad to turn on the transistor. The multilevel switching characteristics of the 1T-1R integrated device were also demonstrated under the DC scheme with the variation of the V_G and V_S , as shown in Fig. S4 (ESI†).

Fig. 6a depicts the potential of the reverse-biased mode for synaptic weight tuning (e.g., neuromorphic computing) applications. The best demonstration of this application came from applying a series of 96 pulses on the 1T-1R integrated device. These pulses simulate the increasing (48 pulses) and decreasing (48 pulses) weights within an artificial neural network (ANN). In particular, a gradual increase in pulse width was applied for both potentiation and depression (20 ns to 490 ns) while keeping the pulse amplitude equal throughout (−2 V and 2.5 V for potentiation and depression, respectively). In total, $>10^6$ increasing and decreasing pulses were conducted on the device in this described method, and this is depicted in Fig. S5a (ESI†) and will be further discussed in the reliability section below. A series of 96 pulses was subjected to the nonlinearity (NL) analysis to investigate the application further quantitatively, in which the data fitted to the following equations:⁴⁸

$$G_{LTP} = G_{\min} + B(1 - e^{-AP}) \quad (3)$$

$$G_{LTD} = G_{\max} + B(1 - e^{A(P-1)}) \quad (4)$$

$$B = \frac{G_{\max} - G_{\min}}{1 - e^{-AP_{\max}}} \quad (5)$$

here, G_{\min} and G_{\max} are the minimum and maximum conductance of the integrated 1T-1R device, and A is the NL factor and is desirable to be as close to 0 as possible, as a perfectly linear PD curve allows maximum compatibility with most neural network models. This leads to a direct mapping of the conductance to weights during the neural network training phase. The inset in Fig. 6a shows that the 1T-1R device achieved an NL factor of 0.45 and 0.14 in PD weight updating with the scheme mentioned above in the reverse-biased mode. This NL analysis was subsequently compared to that operated on the same device in its filamentary mode, and these results are shown in Fig. 6b and c. In these figures, pulse trains of 48 (potentiation) by 48 (depression) pulses were repeated five times, and the data points for each conductance level were presented as min-max error bars. Fig. 6b shows the PD curves for a varied pulse width of 20 ns to 200 ns, keeping the pulse amplitude at 1 V, while Fig. 6c depicts the curves for the varied pulse amplitude of 1 V to 2 V, keeping the pulse width at 20 ns. The NL

factors obtained from operating in the reverse-biased mode were much better than the forward-biased mode ($>>2$). It is also apparent that there is a trade-off between the NL factor and the conductance ratio. A larger pulse width (Fig. 6b) and pulse amplitude (Fig. 6c) generally allow for a larger conductance ratio; however, the NL factor suffers due to the abrupt switching characteristics that filamentary devices inherently possess. This is also exacerbated by the abrupt switching characteristics preventing the full utilization of both the conductance ranges while being able to update the weight (conductance) with each pulse slowly. Additionally, the energy consumption of the resistive switching was also calculated by using $W = V_{\text{peak}} \times I_{\text{peak}} \times t$, where W , V_{peak} , I_{peak} , and t are the energy consumption per switch, peak input voltage, peak output current, and applied pulse width, respectively. The peak voltages and currents were used in the calculation to prevent the underreporting of the power draw. The resultant energy consumption of the device in its reverse-biased mode for synaptic applications was ~ 150 fJ during its first potentiation pulse in the lowest conductance to ~ 95 pJ during its last pulse (48th pulse) in the highest conductance and similarly, from ~ 4 fJ in its first depression pulse to ~ 470 fJ in its final pulse (48th pulse). The device was then investigated for its potential in-memory applications. The integrated 1T-1R device was subjected to alternating positive (2.5 V) and negative (−2.5 V) pulses of 50 ns in its forward-biased mode for 100 pulse cycles. A read pulse of 0.1 V was applied for 100 μ s after every switching pulse to read the conductance levels. The results are depicted in Fig. 6d, where the gate voltage (V_G) varies from 1.9 V to 2.5 V in increments of 0.2 V, demonstrating its capability as an integrated 1T-1R multilevel cell with 4 HCS and 1 LCS. The device's energy consumption to achieve multilevel states was subsequently calculated to be ~ 37 pJ to ~ 60 pJ for the set operation and ~ 40 pJ to ~ 49 pJ for the reset operation. This was compared to the device operating in the reverse-biased mode, where relatively stable binary switching only occurred when applying a minimum pulse of −2.5 V and 2.5 V at 100 μ s for set and reset operations, respectively. The energy consumption of the device operated in both modes was compared in Fig. S6 (ESI†), in which, due to the longer pulse required for the switching, the energy requirements per switch was $\sim 500\times$ lower when operated in the forward-biased mode compared to the reverse-biased mode for the memory scheme. Nevertheless, when comparing the energy consumption of the forward-biased mode memory scheme and the reverse-biased mode weight tuning scheme, the latter requires energy in the same order per switch (~ 10 pJ) in its worst-case scenario (48th pulse during potentiation) as the memory scheme in the forward-biased mode. The demonstrations above show that the device presented in this work is suitable for synaptic tuning applications in the reverse-biased mode and memory applications in the forward-biased mode based on the different requirements of the desired applications.

Additionally, Fig. 6e and f depict the device-to-device variation in both reverse and forward-biased modes using five devices each. In Fig. 6e, the potentiation and depression curves



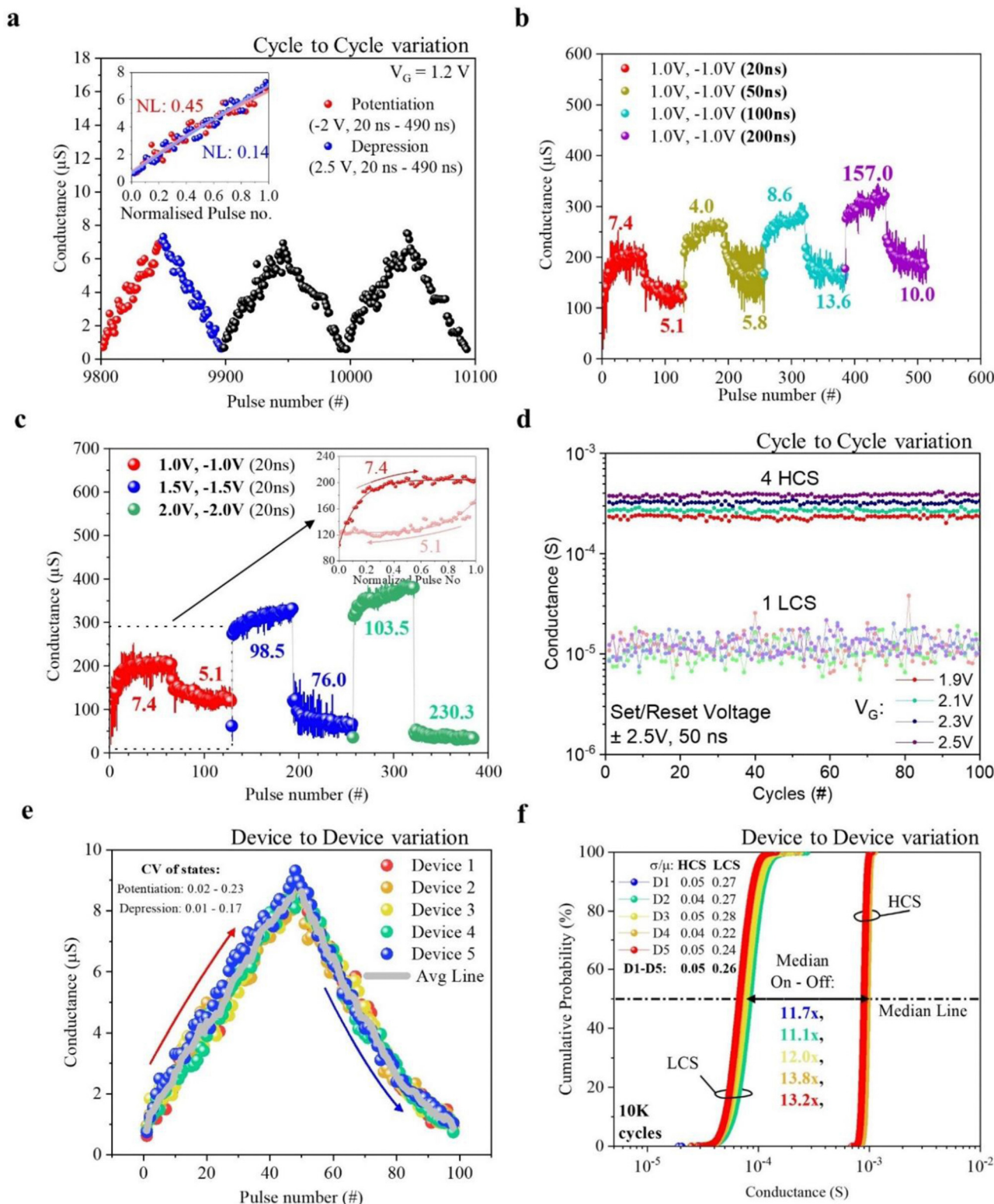


Fig. 6 1T-1R operation of the Pt/Ta₂O₅/Ta₂N device in the reverse-biased mode, and synaptic weight characteristics when applying increasing pulse width from 20 ns to 490 ns. This was compared to operating in the forward-biased mode, where the pulse widths and amplitudes were varied and shown in (b) and (c), respectively. The device was also investigated for its memory applications by operating in the forward-biased mode for binary switching shown in (d), and five states were obtained when varying the V_G from 1.9 V to 2.5 V and switching parameters of 2.5 V for 50 ns. Device to device variation of (e) potentiation and depression in the reverse-biased mode, and (f) binary switching in the forward-biased mode showing low variation between five devices.



of five devices were overlapped while operated in the reverse-biased mode, and the coefficient of variability (CV, or σ/μ) was calculated for each of the 98 states. The CV of the potentiation pulses ranged from 0.02 to 0.23, while the depression pulses ranged from 0.01 to 0.17, showing low deviation of data in each state. The devices were then formed and operated in their forward-biased modes, and all were used to run 10 K cycles each. Fig. 6f shows these 10 K cycles in the form of a cumulative probability plot. Both the LCS and HCS showed low CV, and when comparing inter-device and intra-devices, the CV showed negligible differences. The on-off ratio between the LCS and HCS is also about ~ 12.5 times, meaning that it was still easy to differentiate between the on and off states in the forward-biased mode.

Device reliability

Retention and endurance measurements were conducted to investigate the reliability of the devices discussed in this work. Fig. 7a depicts the retention reliability of the Pt/Ta₂O₅/Ta₂N memristive device structures. The conductance states of a single device were switched (*via* DC measurements, parameters shown in Fig. 7a) at room temperature while being subjected to temperature stress at 85 °C for 10⁴ s for each state. The memristive device was subjected to temperature stress from a pristine condition. It was observed that the conductance state of this pristine device dropped to sub- μ S during this stage. Subsequently, during the reading of the reverse-biased LCS, the device's conductance rose from about 50 s; nevertheless, it reached a stable state above the pristine stressed conductance. Hence, the device was seen to have good retention, and its conductance states remained stable with minimum fluctuations when subjected to temperature stress. The device was also characterized for its endurance, as shown in Fig. 7b, by operating identically to Fig. 6d, whereby 2.5 V and -2.5 V pulses were applied for 50 ns for set and reset operations

(and $V_G = 2.0$ V) in the forward-biased mode. Resultant states were read at 0.1 V for 100 μ s for every cycle. The device was observed to be able to switch for $>10^6$ cycles with stable HCS. However, the LCS tended to fluctuate around 10^{-4} to 10^{-5} S. The endurance in the reverse-biased mode is also shown in Fig. S5a (ESI[†]), by which operating identically to the method to obtain results shown in Fig. 6a, the device continued to work even after $>10^6$ pulses ($>10\,204$ potentiation and depression complete cycles). Figures S5b to S5f (ESI[†]) show the evolution of the PD cycles over the 10^6 pulses, three sets for every order of PD cycles. The NL factor was utilized to quantify the degradation of the device, shown by the fitting in the inset in the aforementioned figures. It was also shown that there were some ~ 1000 pulses that should be run before the device would settle in the optimum state in this application, exhibiting an NL factor of <2 , and after ten cycles, this improved to a value of <1 . Further PD cycles displayed an even more significantly improved NL factor of <0.5 before showing some decline in the potentiation NL factor to >2 after 10^3 PD cycles. Nevertheless, its depression NL factor remained at an excellent level of <0.5 . At 10^4 PD cycles, an overall reduction of the total conductance range caused a more significant degradation in the NL factor of <3 . Additionally, the devices were also shown to have a possibility of reversibility, and the method to achieve this is shown in Fig. S7 (ESI[†]).

Comparison with past works

Table 1 summarizes the comparative analysis of the performance and reliability of devices that exhibit the coexistence of abrupt and gradual switching characteristics presented in this work and those investigated by similar studies in the existing literature. Unlike this work, devices with such duality of modes are not typically explored for practical applications due to the challenges associated with the unintended switching between

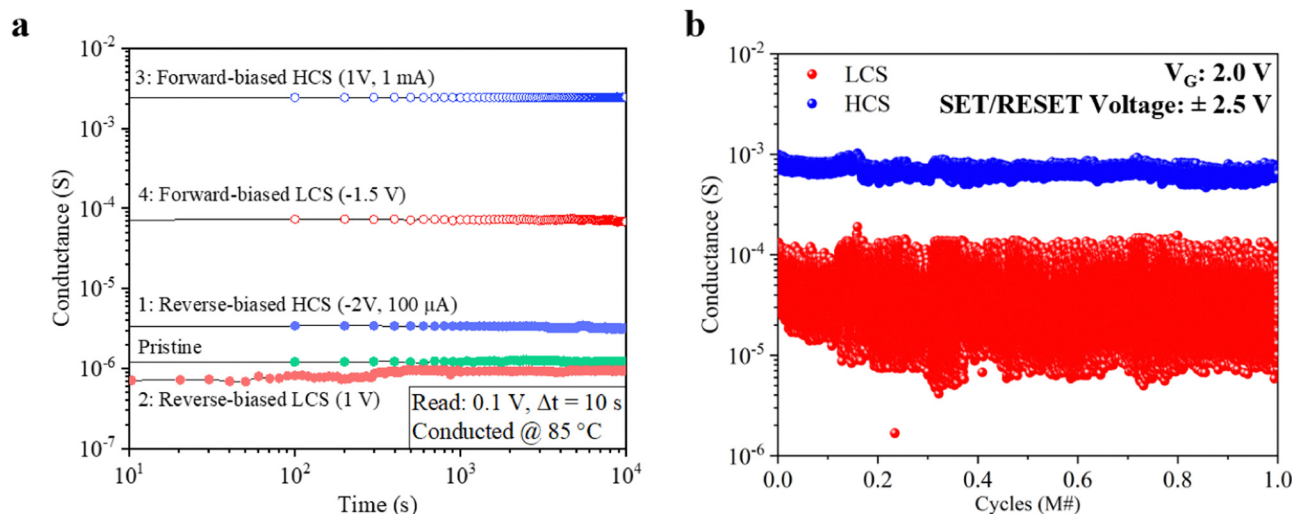


Fig. 7 (a) Retention characteristics of the memristive device show good retention under temperature stress at 85 °C for 10⁴ s. (b) Endurance characterization of switching under 2.5 V and -2.5 V pulses applied for 50 ns for set and reset operations (and $V_G = 2.0$ V) in the forward-biased mode. Resultant states were read at 0.1 V for 100 μ s for every cycle. The device was observed to be able to switch for $>10^6$ cycles with stable LCS and HCS.



Table 1 Comparison of switching performance and reliability of RRAM devices reported in the literature with their abrupt and gradual switching characteristics coexisting within a single device

Memristor structure (thickness [nm])	Device size	Switching performance			Reliability		
		V_{Forming} (V)	$V_{\text{Set}} (V)/V_{\text{Reset}} (V)$ (Switching speed [if avail]) ^c	I_{CC} (mA)	Endurance (cycles)	Retention	Ref.
Au/Sr ₂ TiO ₄ (10)/Nb:SrTiO ₃	100–400 μm	2–3	–4 V/4 V (gradual) 4 V/–10 V (gradual)	10	NA.	NA.	26
Pt(100)/Fe:SrTiO ₃ (500)/ Nb:SrTiO ₃ (substrate)	10 μm	12	–2.2 V/2.8 V (gradual) 2.8 V/–3.5 V (abrupt)	2	NA.	NA.	62
W/graded W _{0.8} (15– 160)/Pt(70)	190–250 nm	–2.2	Pulsed: 1.2 V/–1.6 V (1 μs) (abrupt) –1 V/1.25 V (1 ms) (gradual)	10	2000	10 ⁴ s @ 85 °C ^e	6,24
TiN/DyMn ₂ O ₅ (10)/Pt	N.A.	Yes ^a	1 V/–1.8 V (abrupt) –1.8 V/0.6 V (gradual)	10	100	10 ⁴ s @ 85 °C	63
Pt(100)/ZnO(100)/Pt(100)	50–200 μm	2.8	–2 V/2 V (gradual, bipolar) 1.5 V/0.7 V (abrupt, unipolar)	10 100	100	10 ⁴ s @ 85 °C ^e	64
Pt (20)/Ti(10)/TiO ₂ (3–6)/Pt	60–100 nm	No ^b	–3.5 V/1.7 V (abrupt) 1.7 V/–3.5 V (gradual) Pulsed: –2.1 V (100 ns) (abrupt) 2.2 V (100 μs) (gradual)	0.1	NA.	1.26 × 10 ⁴ s @ 85 °C	28
Pt(30)/Ta ₂ O ₅ (3)/Ta ₂ N	100 nm	1.32	0.7 V/–1 V (abrupt) –2 V/0.8 V (gradual) Pulsed: 2.5 V/–2.5 V (abrupt) –2 V/0.8 V (gradual)	1 0.1 < 0.1 ^d	10 ⁶	10 ⁴ s @ 85 °C	This work

^a The exact forming voltage was not mentioned. ^b Initially conductive. ^c Values reported under DC characterization if not explicitly mentioned.^d I_{CC} imposed with the use of the NMOS transistor. ^e LRS drift under the gradual switching scheme; other states remain stable.

modes. Nevertheless, our proposed memristive device boasts a lower forming voltage (1.32 V) and lower switching voltage (± 2.5 V) relative to the faster switching speed (50 ns). Our device size is also comparably one of the smallest and operates in a lower current range, affirming its potential for further scalability within array integration. Additionally, our devices demonstrate reliability, adhering to industry standards, with stable retention adhering to standards of 10⁴ s at 85 °C. They also exhibit excellent endurance of > 10⁶ set and reset cycles, which was tested with endurance characterization methods recommended by Lanza *et al.*⁶¹ In essence, compared to past studies, this work showcases the robustness of our devices, offering a combination of performance, reliability, and scalability when operated in both forward and reverse-biased modes.

Conclusions

In summary, a Ta₂O₅-based memristive device has been proposed and demonstrated for the coexistence of localized and interfacial switching characteristics within a single device, operated in its forward or reverse-biased mode, respectively. A conductance size-dependence study then revealed the size-independent characteristics of the localized (filamentary) switching states, while interfacial switching states exhibited a size-dependent behavior. Additionally, we proposed and proved that the interfacial switching behavior happened due to the modulation of the oxygen-deficient interfacial layers sandwiched between the active Ta₂N electrode and the Ta₂O₅ oxide

layers. Furthermore, the active electrode (Ta₂N) was chosen due to the reduced oxygen affinity compared to Ta, enabling the filamentary switching within the memristive device. The integration of the 1T–1R device demonstrated its capability as a non-volatile memory element in the forward-biased filamentary mode and a synaptic weight-tuning unit in the reverse-biased non-filamentary mode. The devices showed great endurance of > 10⁶ cycles and pulses in both forward and reverse-biased modes, respectively. The conductance states of the device also exhibited stable states with minimal fluctuations under temperature stress at 85 °C for 10⁴ s. Overall, achieving dual modes within the same device presented in this work unveils the suitability of the device as a unified memory solution.

Methodology

Device fabrication

For the 1T1R integration, a foundry-obtained chip containing NMOS transistor was patterned with electron-beam lithography (EBL) and etched with reactive ion etching (RIE) to access the Ta₂N *via*. A 3 nm-ultrathin Ta₂O₅ switching layer was then deposited by 50 W RF magnetron sputtering at 2 mTorr pressure with 20 sccm Ar gas flow, followed by deposition of a 15 nm Pt top electrode by 50 W DC magnetron sputtering at the same pressure and gas flow.

For the 1R device size variation experiments (Fig. 2a and 5b), a 30 nm Ta₂N was first blanket-deposited on a SiO₂ wafer as a common layer with reactive radio frequency (RF) magnetron



sputtering at 4 mTorr pressure with 26 sccm of Ar and 4 sccm of N₂ gas, resulting in Ta₂N with a sheet resistivity of 270 $\mu\Omega\text{-cm}$ (measured with a four-point technique).⁶⁵ The resultant layer was identical to the Ta₂N from the *via* in the 1T1R chip. Following this, a 30 nm Si₃N₄ passivation layer was deposited using plasma-enhanced chemical vapor deposition (PE-CVD). An EBL step was subsequently done to pattern circular holes that varied from 200 nm to 1 μm . RIE was then used to etch the patterned hole of Si₃N₄ at a rate of 1 nm s⁻¹. The e-beam resist was removed, and another lithography step was done to pattern the top electrode. The 3 nm Ta₂O₅ switching layer and 30 nm Pt TE were then deposited with steps identical to the 1T1R process to fill in the hole which is connected to the Pt.

Materials and device characterisation

For bulk characterization, XPS was conducted by utilizing the Kratos Analytical Axis Supra Plus XPS system.

For surface and topology characterization studies, the non-contact mode of the Park Systems NX1 AFM was utilized to scan the electrode surfaces, and the XRR and XRD measurements were performed using the Rigaku SmartLab with the Cu K α radiation.

The electrical *I-V* and *C-V* characterization studies of the devices were conducted using the Keithley 4200A-SCS semiconductor parameter analyzer.

Author contributions

E. K. K. and P. A. D. conceptualized this work. E. K. K. contributed to most of the work, including research methodology, data curation, formal investigation, and data visualization. H. Y. P. assisted in performing XRD and XRR measurements. P. A. D., H. Y. P., L. L., C. X. X. L., and J. R. T. contributed to the formal analysis of the results. Y. S. Y. and W. S. L. supervised the research. W. S. L. acquired the funds for this research work. E. K. K. drafted the manuscript. All authors reviewed and revised the manuscript.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was supported by a RIE2020 ASTAR AME IAF-ICP Grant (No. I1801E0030). We would also like to acknowledge Dr Teddy Salim from the Facility for Analysis, Characterisation, Testing and Simulation, Nanyang Technological University, Singapore, for the assistance in utilizing the X-ray facilities.

References

- 1 D. Ielmini and H. S. P. Wong, *Nat. Electron.*, 2018, **1**, 333–343.
- 2 W. Zhang, B. Gao, J. Tang, X. Li, W. Wu, H. Qian and H. Wu, *Phys. Status Solidi RRL*, 2019, **13**, 1–13.
- 3 S. C. W. Chow, P. A. Dananjaya, J. M. Ang, D. J. J. Loy, J. R. Thong, S. W. Hoo, E. H. Toh and W. S. Lew, *Appl. Surf. Sci.*, 2023, **608**, 155233.
- 4 P. A. Dananjaya, D. J. J. Loy, S. C. W. Chow and W. S. Lew, *ACS Appl. Electron. Mater.*, 2019, **1**, 2076–2085.
- 5 T. S. Lee, N. J. Lee, H. Abbas, H. H. Lee, T.-S. Yoon and C. J. Kang, *ACS Appl. Electron. Mater.*, 2020, **2**, 1154–1161.
- 6 K. P. Biju, X. Liu, M. Siddik, S. Kim, J. Shin, I. Kim, A. Ignatiev and H. Hwang, *J. Appl. Phys.*, 2011, **110**, 064505.
- 7 M. Kim, M. A. Rehman, D. Lee, Y. Wang, D.-H. Lim, M. F. Khan, H. Choi, Q. Y. Shao, J. Suh, H.-S. Lee and H.-H. Park, *ACS Appl. Mater. Interfaces*, 2022, **14**, 44561–44571.
- 8 H. Ryu and S. Kim, *Nanomaterials*, 2020, **10**, 2055.
- 9 A. Schönhals, C. M. M. Rosário, S. Hoffmann-Eifert, R. Waser, S. Menzel and D. J. Wouters, *Adv. Electron. Mater.*, 2018, **4**, 1700243.
- 10 C. La Torre, A. Kindsmüller, D. J. Wouters, C. E. Graves, G. A. Gibson, J. P. Strachan, R. S. Williams, R. Waser and S. Menzel, *Nanoscale*, 2017, **9**, 14414–14422.
- 11 A. Sebastian, M. Le Gallo, R. Khaddam-Aljameh and E. Eleftheriou, *Nat. Nanotechnol.*, 2020, **15**, 529–544.
- 12 S. Yu, W. Shim, X. Peng and Y. Luo, *IEEE Trans. Circuits Syst. Regul. Pap.*, 2021, **68**, 2753–2765.
- 13 S. Cho, J. Jung, S. Kim and J. J. Pak, *Semicond. Sci. Technol.*, 2020, **35**, 085006.
- 14 L. Liu, P. A. Dananjaya, C. C. I. Ang, E. K. Koh, G. J. Lim, H. Y. Poh, M. Y. Chee, C. X. X. Lee and W. S. Lew, *Nanoscale*, 2023, **15**, 17076–17084.
- 15 L. Liu, P. A. Dananjaya, M. Y. Chee, G. J. Lim, C. X. X. Lee and W. S. Lew, *ACS Appl. Mater. Interfaces*, 2023, **15**, 29287–29296.
- 16 M. Prezioso, F. Merrih-Bayat, B. D. Hoskins, G. C. Adam, K. K. Likharev and D. B. Strukov, *Nature*, 2015, **521**, 61–64.
- 17 C. Li, D. Belkin, Y. Li, P. Yan, M. Hu, N. Ge, H. Jiang, E. Montgomery, P. Lin, Z. Wang, W. Song, J. P. Strachan, M. Barnell, Q. Wu, R. S. Williams, J. J. Yang and Q. Xia, *Nat. Commun.*, 2018, **9**, 2385.
- 18 M. Zhao, B. Gao, J. Tang, H. Qian and H. Wu, *Appl. Phys. Rev.*, 2020, **7**, 011301.
- 19 W. Wan, R. Kubendran, C. Schaefer, S. B. Eryilmaz, W. Zhang, D. Wu, S. Deiss, P. Raina, H. Qian, B. Gao, S. Joshi, H. Wu, H.-S. P. Wong and G. Cauwenberghs, *Nature*, 2022, **608**, 504–512.
- 20 W.-H. Chen, C. Dou, K.-X. Li, W.-Y. Lin, P.-Y. Li, J.-H. Huang, J.-H. Wang, W.-C. Wei, C.-X. Xue, Y.-C. Chiu, Y.-C. King, C.-J. Lin, R.-S. Liu, C.-C. Hsieh, K.-T. Tang, J. J. Yang, M.-S. Ho and M.-F. Chang, *Nat. Electron.*, 2019, **2**, 420–428.
- 21 B. Tang, H. Veluri, Y. Li, Z. G. Yu, M. Waqar, J. F. Leong, M. Sivan, E. Zamburg, Y.-W. Zhang, J. Wang and A. V. Y. Thean, *Nat. Commun.*, 2022, **13**, 3037.
- 22 J. Jian, P. Dong, Z. Jian, T. Zhao, C. Miao, H. Chang, J. Chen, Y.-F. Chen, Y.-B. Chen, H. Feng and B. Sorli, *ACS Nano*, 2022, **16**, 20445–20456.
- 23 R. Muenstermann, T. Menke, R. Dittmann and R. Waser, *Adv. Mater.*, 2010, **22**, 4819–4822.



- 24 K. P. Biju, X. Liu, S. Kim, S. Jung, J. Park and H. Hwang, *Phys. Status Solidi RRL*, 2011, **5**, 89–91.
- 25 S. Yun, J. Park, M. Kang and S. Kim, *Results Phys.*, 2022, **34**, 105307.
- 26 K. Shibuya, R. Dittmann, S. Mi and R. Waser, *Adv. Mater.*, 2010, **22**, 411–414.
- 27 F. Miao, J. Joshua Yang, J. Borghetti, G. Medeiros-Ribeiro and R. Stanley Williams, *Nanotechnology*, 2011, **22**, 254007.
- 28 H. Zhang, S. Yoo, S. Menzel, C. Funck, F. Cüppers, D. J. Wouters, C. S. Hwang, R. Waser and S. Hoffmann-Eifert, *ACS Appl. Mater. Interfaces*, 2018, **10**, 29766–29778.
- 29 M. Ismail, H. Abbas, A. Sokolov, C. Mahata, C. Choi and S. Kim, *Ceram. Int.*, 2021, **47**, 30764–30776.
- 30 K. X. Shi, H. Y. Xu, Z. Q. Wang, X. N. Zhao, W. Z. Liu, J. G. Ma and Y. C. Liu, *Appl. Phys. Lett.*, 2017, **111**, 223505.
- 31 R. Simpson, R. G. White, J. F. Watts and M. A. Baker, *Appl. Surf. Sci.*, 2017, **405**, 79–87.
- 32 S. Khan, M. J. M. Zapata, M. B. Pereira, R. V. Gonçalves, L. Strizik, J. Dupont, M. J. L. Santos and S. R. Teixeira, *Phys. Chem. Chem. Phys.*, 2015, **17**, 23952–23962.
- 33 D. Cristea, L. Cunha, C. Gabor, I. Ghiuta, C. Croitoru, A. Marin, L. Velicu, A. Besleaga and B. Vasile, *Nanomaterials*, 2019, **9**, 476.
- 34 Q. Zhou and J. Zhai, *Phys. Status Solidi A*, 2014, **211**, 173–179.
- 35 M. Ismail, E. Ahmed, A. M. Rana, I. Talib, T. Khan, K. Iqbal and M. Y. Nadeem, *Thin Solid Films*, 2015, **583**, 95–101.
- 36 A. Wedig, M. Luebben, D.-Y. Cho, M. Moors, K. Skaja, V. Rana, T. Hasegawa, K. K. Adepalli, B. Yildiz, R. Waser and I. Valov, *Nat. Nanotechnol.*, 2016, **11**, 67–74.
- 37 K. Baek, S. Park, J. Park, Y.-M. Kim, H. Hwang and S. H. Oh, *Nanoscale*, 2017, **9**, 582–593.
- 38 P. Huang, X. Y. Liu, W. H. Li, Y. X. Deng, B. Chen, Y. Lu, B. Gao, L. Zeng, K. L. Wei, G. Du, X. Zhang and J. F. Kang, *Int. Electron Devices Meet.*, 2012, 26.6.1–26.6.4.
- 39 Y. D. Zhao, P. Huang, Z. Chen, C. Liu, H. T. Li, B. Chen, W. J. Ma, F. F. Zhang, B. Gao, X. Y. Liu and J. F. Kang, in 2015 International Symposium on VLSI Technology, Systems and Applications, 2015, 1–2.
- 40 C. Cai, S. Wei, Z. Yin, J. Bai, W. Xie, Y. Li, F. Qin, Y. Su and D. Wang, *Appl. Surf. Sci.*, 2021, **560**, 149960.
- 41 B. Widrow, Y. Kim, D. Park and J. K. Perin, *Artificial Intelligence in the Age of Neural Networks and Brain Computing*, ed. R. Kozma, C. Alippi, Y. Choe and F. C. Morabito, Academic Press, 2019, pp. 1–30.
- 42 F.-C. Chiu, *Adv. Mater. Sci. Eng.*, 2014, **2014**, 578168.
- 43 W. Banerjee, *Electronics*, 2020, **9**, 1029, DOI: [10.3390/electronics9061029](https://doi.org/10.3390/electronics9061029).
- 44 A. Lekshmi Jagath, C. Hock Leong, T. N. Kumar and H. F. Almurib, *J. Eng.*, 2019, **2019**, 4644–4652.
- 45 J. Lin, N. Masaaki, A. Tsukune and M. Yamada, *Appl. Phys. Lett.*, 1999, **74**, 2370–2372.
- 46 D. Cristea, A. Crisan, N. Cretu, J. Borges, C. Lopes, L. Cunha, V. Ion, M. Dinescu, N. P. Barradas, E. Alves, M. Apreutesei and D. Munteanu, *Appl. Surf. Sci.*, 2015, **354**, 298–305.
- 47 C. S. Kang, H.-J. Cho, Y. H. Kim, R. Choi, K. Onishi, A. Shahriar and J. C. Lee, *J. Vac. Sci. Technol., B: Microelectron. Nanometer Struct.–Process., Meas., Phenom.*, 2003, **21**, 2026–2028.
- 48 *CRC handbook of chemistry and physics*, CRC Press, 103th edn, Rumble, J. R., Ed.; CRC Press, 2022.
- 49 X. Li, H. Wu, B. Gao, W. Wu, D. Wu, N. Deng, J. Cai and H. Qian, *Nanotechnology*, 2016, **27**, 305201.
- 50 L. Goux, A. Fantini, Y. Y. Chen, A. Redolfi, R. Degraeve and M. Jurczak, *ECS Solid State Lett.*, 2014, **3**, Q79.
- 51 V. Y.-Q. Zhuo, Y. Jiang, M. H. Li, E. K. Chua, Z. Zhang, J. S. Pan, R. Zhao, L. P. Shi, T. C. Chong and J. Robertson, *Appl. Phys. Lett.*, 2013, **102**, 062106.
- 52 L. Goux, A. Fantini, A. Redolfi, C. Y. Chen, F. F. Shi, R. Degraeve, Y. Y. Chen, T. Witters, G. Groeseneken and M. Jurczak, in 2014 Symposium on VLSI Technology (VLSI-Technology): Digest of Technical Papers, 2014, pp. 1–2.
- 53 C. Sun, S. M. Lu, F. Jin, W. Q. Mo, J. L. Song and K. F. Dong, *J. Alloys Compd.*, 2018, **749**, 481–486.
- 54 A. Jiang, T. A. Tyson, L. Axe, L. Gladczuk, M. Sosnowski and P. Cote, *Thin Solid Films*, 2004, **479**, 166–173.
- 55 A. Zaman and E. I. Meletis, *Coatings*, 2017, **7**, 209.
- 56 K. C. Chiang, C. H. Cheng, H. C. Pan, C. N. Hsiao, C. P. Chou, A. Chin and H. L. Hwang, *IEEE Electron Device Lett.*, 2007, **28**, 235–237.
- 57 C. Wenger, M. Lukosius, H.-J. Müssig, G. Ruhl, S. Pasko and C. Lohe, *J. Vac. Sci. Technol., B: Microelectron. Nanometer Struct.–Process., Meas., Phenom.*, 2009, **27**, 286–289.
- 58 C. H. Cheng, H. H. Hsu, W. B. Chen, A. Chin and F. S. Yeh, *Electrochem. Solid-State Lett.*, 2009, **13**, H16.
- 59 T. Kim, H. Son, I. Kim, J. Kim, S. Lee, J. K. Park, J. Y. Kwak, J. Park and Y. Jeong, *Sci. Rep.*, 2020, **10**, 11247.
- 60 A. Schonhals, A. Kindsmuller, C. L. Torre, H. Zhang, S. Hoffmann-Eifert, S. Menzel, R. Waser and D. J. Wouters, *IEEE International Memory Workshop*, 2017, 1–4.
- 61 M. Lanza, R. Waser, D. Ielmini, J. J. Yang, L. Goux, J. Suñe, A. J. Kenyon, A. Mehonic, S. Spiga, V. Rana, S. Wiefels, S. Menzel, I. Valov, M. A. Villena, E. Miranda, X. Jing, F. Campabadal, M. B. Gonzalez, F. Aguirre, F. Palumbo, K. Zhu, J. B. Roldan, F. M. Puglisi, L. Larcher, T.-H. Hou, T. Prodromakis, Y. Yang, P. Huang, T. Wan, Y. Chai, K. L. Pey, N. Raghavan, S. Dueñas, T. Wang, Q. Xia and S. Pazos, *ACS Nano*, 2021, **15**, 17214–17231.
- 62 D. Cooper, C. Baeumer, N. Bernier, A. Marchewka, C. La Torre, R. E. Dunin-Borkowski, S. Menzel, R. Waser and R. Dittmann, *Adv. Mater.*, 2017, **29**, 1700212.
- 63 Y.-T. Tsai, T.-C. Chang, W.-L. Huang, C.-W. Huang, Y.-E. Syu, S.-C. Chen, S. M. Sze, M.-J. Tsai and T.-Y. Tseng, *Appl. Phys. Lett.*, 2011, **99**, 092106.
- 64 C.-H. Huang, J.-S. Huang, C.-C. Lai, H.-W. Huang, S.-J. Lin and Y.-L. Chueh, *ACS Appl. Mater. Interfaces*, 2013, **5**, 6017–6023.
- 65 H. Y. Poh, C. C. I. Ang, T. L. Jin, F. N. Tan, G. J. Lim, S. Wu, F. Poh and W. S. Lew, *Appl. Phys. Lett.*, 2022, **121**, 012405.

