

DS 34: Resistive Switching (jointly with DF, KR, HL)

Time: Friday 9:30–12:45

Location: H32

DS 34.1 Fri 9:30 H32

Ab initio study of defects in SrTiO₃ bulk and (100) surfaces

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Oxygen vacancies are believed to play a major role in the conduction mechanisms that enable resistive switching in oxide materials. Employing density functional theory (DFT) and the DFT+U model, we use the full-potential linearized augmented plane wave method as implemented in the FLEUR code to study the formation of point defects in the perovskite SrTiO₃ with varying coordination. We calculated the formation energy of an O-vacancy in both bulk supercells and (100) surface including different, c(2×2) and p(2×2), in-plane unit cells and different terminations. After performing full relaxation, we found that the bulk and SrO-terminated surface have a nonmagnetic, while TiO₂-terminated surface has a ferromagnetic solution. Using the c(2×2) unit cell, the vacancy formation energy was smaller for the bulk than for the SrO- and even TiO₂-terminated surface. On the other hand, the p(2×2) unit cell shows that TiO₂-terminated surface has the lowest formation energy, more than 1 eV lower than the bulk value. Similar comparisons will be presented when including the DFT+U model that is used to correct the bulk bandgap and improve the localization of the defect states.

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DS 34.2 Fri 9:45 H32

Resistive switching properties in ion beam modified SrTiO₃

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Resistive switching phenomena, which are for instance observed in perovskite-type transition metal oxides, attract intensive attention for their potential application in future nonvolatile memory. Strontium titanate (SrTiO₃) exhibits bipolar resistive switching between a high- and a low-resistance state when applying an appropriate electric field. It is often proposed that the underlying mechanism for bipolar resistive switching in SrTiO₃ originates from oxygen-vacancy migration along filaments based on extended defects such as dislocations or grain boundaries.

Here we report on well-defined damage formation due to ion irradiation which allows a better control of the lateral and vertical defect arrangements and concentrations. Therefore, we deposited 100 nm single crystalline SrTiO₃ thin films with low intrinsic defect concentration on niobium doped SrTiO₃ substrates by pulsed laser deposition and implanted these samples with swift heavy gold ions. After irradiation the films were characterized using transmission electron microscopy and Rutherford backscattering spectrometry. Under ion irradiation, the as-deposited crystalline films undergo amorphisation due to the formation and overlap of amorphous tracks. The electrical properties of SrTiO₃, i.e. the resistive switching properties are discussed in terms of damage concentration.

DS 34.3 Fri 10:00 H32

Cation defect engineering in SrTiO₃ thin films by PLD with Verification and implication on memristive properties— SEBASTIAN WICKLEIN¹, ●CHENCHENG XU¹, ALESSIA SAMBRI², SALVATORE AMORUSO², DAVID KEEBLE³, ANNEMARIE KÖHL¹, WERNER EGGER⁴, and REGINA DITTMANN¹ — ¹Peter Grünberg Institut 7, Forschungszentrum Jülich GmbH, Germany — ²Università degli Studi di Napoli Federico II, Dipartimento di Scienze Fisiche & CNR-SPIN, I-80126 Napoli, Italy — ³University of Dundee, School of Engineering, Physics and Mathematics, Dundee DD1 4HN, Scotland — ⁴University Bundeswehr, D-85577 Munich, Germany

The origin of the c-axis expansion in homoepitaxial STO thin films is investigated by positron annihilation lifetime spectroscopy (PALS): Low laser fluence results in Ti vacancy rich sample while high laser fluence for the Sr vacancy rich sample.

XPS measurement on the ablated spot on the targets shows that increased laser fluence ablates more Ti. The ToF (Time of Flight) data from OES (optical emission spectrometry) indicate a preferred scattering of Ti because of background gas. The two effects together

lead to tunable stoichiometry of the film.

In the MIM (metal insulator metal) structure Sr-rich films exhibit the most stable switching behavior and highest on/off ratio, while in the LC AFM (local conducting atomic force microscopy) switching the on/off ratio of Ti is the highest.

DS 34.4 Fri 10:15 H32

Resistive Switching in thermally oxidized Titanium— ●DANIEL BLASCHKE¹, ILONA SKORUPA¹, BERND SCHEUMANN¹, ANDREA SCHOLZ¹, PETER ZAHN¹, SIBYLLE GEMMING¹, KAY POTZGER¹, AGNIESZKA BOGUSZ², and HEIDEMARIE SCHMIDT² — ¹Helmholtz-Zentrum Dresden-Rossendorf, Institute of Ion Beam Physics and Materials Research, P.O. Box 510119, 01314 Dresden - Germany — ²Dept. Electr. Eng. & Inf. Techn., TU Chemnitz, 09107 Chemnitz

In recent years the resistive switching of binary transition metal oxides like NiO, Nb₂O₅ and TiO₂ has attracted considerable attention for application in nonvolatile memory storage systems.

For our investigations we used a thin rutile TiO₂ film, which was prepared by the thermal oxidation of a 100nm thick e-beam evaporated Ti film. The oxidation temperatures were varied from 500°C to 800°C at an oxygen partial pressure of 1 atmosphere. We will present the dependence of the crystal structure and the switching behavior on the oxidation temperature as well as an interesting feature on the time-dependent evolution of the resistance during the Reset process.

The project is funded by the Initiative and Networking Fund of the Helmholtz Association (VH-VI-422).

DS 34.5 Fri 10:30 H32

Non-volatile resistive switching in multiferroic YMnO₃ thin films— ●AGNIESZKA BOGUSZ^{1,2}, ILONA SKORUPA¹, ANDREA SCHOLZ¹, OLIVER G. SCHMIDT^{2,3}, and HEIDEMARIE SCHMIDT² — ¹Institute of Ion Beam Physics and Materials Research, Helmholtz-Zentrum Dresden-Rossendorf, 01314 Dresden, Germany — ²Faculty of Electrical Engineering and Information technology, Chemnitz University of Technology, 09107 Chemnitz, Germany — ³Institute for Integrative Nanosciences, IFW-Dresden, 01069 Dresden, Germany

Intensive research on multiferroic materials [1] is driven by the possibility of creating novel, miniaturized tunable multifunctional devices [2]. This work investigates resistive switching behavior of YMnO₃ thin films, which can be utilized in new generation memory devices. Series of YMnO₃ films were grown by pulsed laser deposition on Si substrates with Pt bottom electrode at temperatures varying between 500°C and 850°C. Characterization of as-grown samples by X-ray diffraction and scanning electron microscopy was followed by determination of electrical properties of films in metal-insulator-metal (MIM) configuration. Results showed that the YMnO₃ films grown at 800°C exhibit the best resistive switching properties with high resistance ratio (>10000) of high over low resistance state. Switching mechanism is ascribed to the structural transitions within the film upon applied current.

[1] A. Bogusz et al., Defect Diffus. Forum 323-325, 115 (2012)

[2] Y. Shuai, H. Schmidt et al., J. Appl. Phys. 109, 124117 (2011); J. Appl. Phys. 111, 07D906 (2012)

DS 34.6 Fri 10:45 H32

Practical guide for validated memristance measurements— ●NAN DU^{1,2}, YAO SHUAI³, WENBO LUO³, CHRISTIAN MAYR⁴, RENE SCHÜFFNY⁴, OLIVER G. SCHMIDT^{1,2}, and HEIDEMARIE SCHMIDT¹ — ¹TU Chemnitz, Faculty of Electrical Engineering and Information Technology, 09107 Chemnitz, Germany — ²Institute for Integrative Nanosciences, IFW Dresden, Helmholtzstraße 20, 01069 Dresden, Germany — ³Helmholtz Research Center Dresden-Rossendorf, 01328 Dresden, Germany — ⁴TU Dresden, Department of Electrical Engineering and Information Technology, 01062 Dresden, Germany

L.O. Chua predicted rather simple charge-flux curves for active and passive memristors and presented active memristor circuit realizations already in the 70s. However, despite the fact that memristors give rise to complicated hysteretic current-voltage curves, memristors are traced in current-voltage curves. Here we give a practical guide how to use normalized charge-flux curves for the prediction of current-voltage characteristics of memristors with stable electrical characteristics in dependence on the shape and amplitude of the input voltage or input current signals. In the case of memristive BiFeO₃ thin film capacitor

structures [1] the normalized charge-flux curves superimpose for different numbers of measurement points and a different measurement time per measurement point. Such normalized charge-flux curve can be used for the prediction of current-voltage characteristics of plastic synapses in neuromorphic systems [2]. [1] Y. Shuai et al., *J. of Appl. Phys.* 109, 124117-124117-4 (2011). [2] C. Mayr et al., NIPS 2012, in press.

Coffee break (15 min)

DS 34.7 Fri 11:15 H32

Creating an Oxygen Gradient in Nb₂O₅ by Argon Irradiation for Resistive Switching Memory — ●HELGE WYLEZICH¹, HANNES MÄHNE¹, DANIEL BLASCHKE², STEFAN SLESAZECK¹, and THOMAS MIKOLJIACK¹ — ¹NamLab gGmbH, Nöthnitzer Str. 64, D-01187 Dresden — ²Helmholtz-Zentrum Dresden-Rossendorf, D-01314 Dresden

It is common knowledge that an oxygen gradient is mandatory for bipolar resistive switching [1]. We confirmed this by investigations of thin films with Nb₂O₅ as switching layer. Samples with two inert Pt electrodes are nearly symmetric and do not show bipolar resistive switching behavior. Replacing one Pt electrode with a reactive one – for example Al or Nb – results in an unsymmetrical device. These samples could be switched reproducibly. It is also possible to create an oxygen gradient by depositing a stack of two different niobium oxide layers. While the first layer consists of stoichiometric Nb₂O₅ the second layer is sputtered substoichiometric [2].

A new approach is to get an oxygen gradient by irradiating the oxide layer with argon. Two effects appear: The argon sputters the surface of the Nb₂O₅ layer and so the oxide thickness decreases. Because the Nb-atoms are heavier than the O-atoms, the oxygen sputter rate is higher and the surface becomes niobium rich. The investigated samples consist of a Pt-Nb₂O₅-Pt stack. The oxide layer was irradiated by different Ar-doses before top electrode deposition. At the highest dose $\Phi = 3e16 \text{ cm}^{-2}$ the resulting oxygen gradient enables resistive switching.

[1] Bertaud et al. (TSF 520, 2012)

[2] Mähne et al. (MEMCOM Workshop 2012)

DS 34.8 Fri 11:30 H32

Multilevel resistive switching in Ar⁺ irradiated BiFeO₃ thin films — ●YAO SHUAI¹, XIN OU², WENBO LUO², NAN DU³, DANILO BÜRGER^{2,3}, OLIVER G. SCHMIDT^{3,4}, and HEIDEMARIE SCHMIDT² — ¹State Key Laboratory of Electronic Thin Films and Integrated Devices, UESTC, China — ²Helmholtz-Zentrum Dresden-Rossendorf e.V., Institute of Ion Beam Physics and Materials Research, Germany — ³University of Technology Chemnitz, Faculty of Electrical Engineering and Information Technology, 09107 Chemnitz, Germany — ⁴Institute for Integrative Nanosciences, IFW Dresden, Helmholtzstraße 20, 01069 Dresden, Germany

Low energy Ar⁺ ion irradiation has been applied to an Au/BiFeO₃/Pt capacitor structures before deposition of the Au top electrode. The irradiated thin films exhibit multilevel resistive switching without detrimental resistance degradation, which makes the intermediate resistance states more distinguishable as compared to the non-irradiated thin film [1]. The stabilization of resistance states after irradiation is discussed based on the analysis of conduction mechanism during the resistive switching in BiFeO₃ with a rectifying Au top electrode and a nonrectifying Pt bottom electrode [2]. Furthermore, it is shown how the conduction mechanisms change from room temperature to 423 K. [1] Y. Shuai, X. Ou et al., *IEEE Device Letters*, 2012, in press. [2] Y. Shuai, S. Zhou, D. Bürger, M. Helm, H. Schmidt, *J. Appl. Phys.* 109 (2011), 124117-4.

DS 34.9 Fri 11:45 H32

Influence of thickness ratio on resistive switching in BiFeO₃:Ti/BiFeO₃ bilayer structures — ●TIANGUI YOU¹, WENBO LUO², YAO SHUAI^{1,2}, NAN DU¹, DANILO BÜRGER^{1,3}, ILONA SKORUPA³, OLIVER G. SCHMIDT^{1,4}, and HEIDEMARIE SCHMIDT¹ — ¹Chemnitz University of Technology, 09107 Chemnitz, Germany — ²University of Electronic Science and Technology of China, 610054 Chengdu, China — ³Helmholtz-Zentrum Dresden-Rossendorf, P.O. Box 510119, 01314 Dresden, Germany — ⁴IFW-Dresden, 01069 Dresden, Germany

Nonvolatile resistive switching in BiFeO₃ (BFO) [1] has attracted increasing attention; however, the underlying resistive switching mech-

anism is still controversial which restricts its application in non-volatile memory devices. BFO:Ti/BFO bilayer structures with a 540 nm thick BFO layer and different thickness of BFO:Ti layer were grown on Pt/Sapphire substrates by pulsed laser deposition using the same growth conditions. Circular Au top electrodes were prepared with magnetron sputtering. Au/BFO/Pt single layer structures show a symmetric I-V curve without hysteresis due to the formation of Schottky contacts at both the top and bottom interface. However, Au/BFO/BFO:Ti/Pt bilayer structures exhibit an obvious resistive switching behavior under both voltage polarities. The influence of the thickness of BFO:Ti on the conduction mechanisms in Au/BFO/BFO:Ti/Pt bilayer structures is discussed to reveal similarities and differences between single and bilayer structures.

Reference [1] Y. Shuai et al., *J. Appl. Phys.*, 109, 124117(2011)

DS 34.10 Fri 12:00 H32

Nanoscale resistive switching in epitaxial and polycrystalline BiFeO₃ thin films — ●YAO SHUAI¹, WENBO LUO¹, CHUANGUI WU¹, WANLI ZHANG¹, OLIVER G. SCHMIDT^{2,3}, and HEIDEMARIE SCHMIDT² — ¹State Key Laboratory of Electronic Thin Films and Integrated Devices, UESTC, China — ²University of Technology Chemnitz, Faculty of Electrical Engineering and Information Technology, 09107 Chemnitz, Germany — ³Institute for Integrative Nanosciences, IFW Dresden, 01069 Dresden, Germany

Nonvolatile [1], bipolar, and multilevel [2] resistive switching has been observed in ca. 500 nm thick polycrystalline BiFeO₃ thin films with rectifying, circular Au top electrodes and a nonrectifying Pt bottom electrode. The diameter of the Au top electrodes amounts to ca. 0.5 μm. By scanning a positionable top contact with a diameter of only 10 nm over polycrystalline BiFeO₃ thin films under a constant applied dc voltage, the high and low resistance state can be locally written and afterwards read. It has been observed that for thinner polycrystalline BiFeO₃ films with a thickness below 300 nm, no resistive switching can be observed either with large or with small scale top contacts. Bipolar resistive switching can also be realized in ca. 50 nm thick epitaxial BiFeO₃ films on SrRuO₃/SrTiO₃ with a positionable top contact. This resistance is mainly determined by the ferroelectric polarization and the barrier height of the top and bottom contact. For thicker epitaxial BiFeO₃ films the unique relation between ferroelectric polarization and resistance state is diminished. [1] Y. Shuai et al., *J. Appl. Phys.* 109 (2011). [2] Y. Shuai et al., *IEEE Device Letters* (2012) in press.

DS 34.11 Fri 12:15 H32

An electronic implementation of amoeba anticipation — ●MIRKO HANSEN¹, KARLHEINZ OCHS², MARTIN ZIEGLER¹, and HERMANN KOHLSTEDT¹ — ¹Faculty of Engineering, Christian-Albrechts-Universität zu Kiel, 24143 Kiel, Germany — ²Ruhr-Universität Bochum, 44780 Bochum, Germany

In nature, the capability to memorize environmental changes can already be observed in unicellular organisms like amoebas[1]. An amoeba changes its locomotive speed when it is exposed to unfavorable conditions. If a series of unfavorable conditions is applied, the amoeba later on behaves similarly on a single incident. Pershin et al.[2] are able to emulate this behavior using a simple resistive switching circuit model consisting of an inductor, a capacitor and a resistive switching device. We experimentally implement this model using a resistive switching device. A theoretical analysis of the circuit is presented to gain further insight into the functionality of this model and to give advice for the implementation of resistive switching devices in LC-circuits.

[1] T. Saigusa, A. Tero, T. Nakagaki, Y. Kuramoto, *Phys. Rev. Lett.* **100**, (2008) 018101

[2] Y. V. Pershin, S. La Fontaine, M. Di Ventra, *Phys. Rev. E* **80**, (2009) 021926

DS 34.12 Fri 12:30 H32

Lattice dynamics in Sb- and Te-based phase-change materials — ●RONNIE ERNST SIMON^{1,2}, ILYA SERGUEEV³, and RAPHAËL PIERRE HERMANN^{1,2} — ¹Jülich Centre for Neutron Science JCNS and Peter Grünberg Institut PGI, JARA-FIT Forschungszentrum Jülich GmbH, D-52425 Jülich, Germany — ²Faculté des Sciences, Université de Liège, B-4000 Liège, Belgium — ³Deutsches Elektronen-Synchrotron, D-22607 Hamburg, Germany

Phase-change materials exhibit a significant change of the optical reflectivity and electrical resistivity upon crystallization which renders these materials applicable for optical storage devices and non-volatile electronic memories. In order to understand the switching kinetics between the amorphous and the metastable crystalline states a detailed

knowledge of the lattice dynamics of the different phases is crucial. A suitable technique for the investigation of lattice dynamics is nuclear inelastic scattering (NIS) which gives access to the element specific density of phonon states (DPS). We performed NIS measurements in Sb- and Te-based phase-change materials in the amorphous and crystalline

phases. We have recently extended the experimental possibilities by demonstrating the feasibility of high pressure NIS measurements, up to 75 GPa, in Sb_2Te_3 . The ESRF is acknowledged for the provision of synchrotron radiation beamtime at ID18.