Interfacial Chemistry, Defects, and Strain in Multiferroic Heterostructures

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The study of multiferroics is a burgeoning field of inquiry [1], partially owing to the hope of exploiting the magnetoelectric effect to permit rapid control of magnetic domain orientation via an applied electric field. Extrinsic multiferroic devices can be fabricated by layering ferroelectric and ferromagnet materials [2] whereby magnetoelectric coupling occurs via a strain-mediated mechanism. Magnetic properties of the structure are also dependent on strain. [3] Relationships between local chemistry, strain, and dislocation cores have been reported. [4] An improved understanding of strain-dependent coupling, interfacial chemistry, and misfit dislocations at the ferroelectric/ferromagnetic interface may allow for the development of models that predict device behavior and failure under an applied electric field.

In these studies, thin films of the ferromagnetic perovskite La_{0.67}Sr_{0.33}MnO (LSMO) are deposited on thin films of the ferroelectric perovskite Pb_{0.20}Zr_{0.80}TiO₃ (PZT) via pulsed laser deposition (PLD). Figure 1 shows the deposition sequence of the structures. Strontium titanate (SrTiO₃) in the (100) orientation serve as the substrate and SrRuO₃ is deposited as a conducting bottom electrode. The thickness of the PZT is varied between 20nm and 50nm while the thickness of the LSMO is constant. Top and bottom conducting electrodes allow for complementary *in-situ* biasing studies. [5] A HR-TEM image showing the LSMO/PZT and PZT/SRO interfaces is shown in Figure 2. The lattice constant of the PZT at the PZT/LSMO interface depends on the PZT thickness, thus the interfacial strain and the strain in the PZT and LSMO layers is varied by controlling the thickness of PZT.

Aberration-corrected, atomically-resolved STEM-EELS and Z-contrast imaging are applied to characterize the local chemistry and defect structure at the ferromagnetic/ferroelectric interface. Strain plays a vital role in determining the concentration and type of interfacial defects and is a crucial variable in defining the magnetoelectric coupling and exchange bias in the system as well as the nucleation, propagation, growth, and hysteresis behavior of ferromagnetic domains.

The ability to control magnetic orientation via magnetoelectric coupling may be doomed by high concentrations of misfit dislocations at the ferromagnetic/ferroelectric interface. Such an interface is anticipated to have a non-uniform electric field distribution. STEM-EELS allows for the electronic and chemical structure of dislocation cores to be studied at the atomic level and makes it possible to measure the diffusion of specific atoms in and around a dislocation core. The question of where oxygen atoms sit in relation to a dislocation core and how cations and oxygen vacancies distribute

may ultimately determine the device properties. A better understanding of the phenomena and structure at the interface between ferroelectric and ferromagnetic materials is reported.

References

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| Au | Electrode |
|--|---------------|
| La _{0.67} Sr _{0.33} MnO ₃ | Ferromagnetic |
| Pb _{0.2} Zr _{0.8} TiO ₃ | Ferroelectric |
| SRO | Electrode |
| STO (100) | Substrate |

Figure 1: Schematic of multiferroic thin film structure fabricated by pulsed laser deposition on STO (100) substrate.

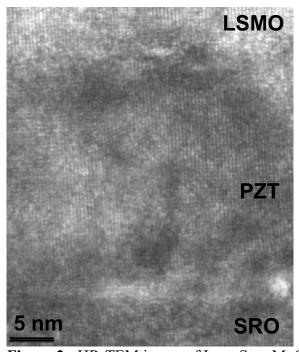


Figure 2: HR-TEM image of La_{0.67}Sr_{0.33}MnO (LSMO) deposited on Pb_{0.20}Zr_{0.80}TiO₃ (PZT).