Phonon confinement and interface lattice dynamics of ultrathin high-\(k\) rare earth sesquioxide films: the case of Eu\(_2\)O\(_3\) on YSZ(001)†

Svetoslav Stankov, \(\ddagger\)*ab Dániel G. Merkel, \(\ddagger\)cd Jochen Kalt, \(\ddagger\)ab Jörg Göttlicher,\(\ddagger\) Jan Łazewski, \(\ddagger\)e Małgorzata Sternik, \(\ddagger\)e Paweł T. Jochym, \(\ddagger\)e Przemysław Piekarz, \(\ddagger\)e Tilo Baumbach, \(\ddagger\)ab Aleksandr I. Chumakov\(\ddagger\) and Rudolf Rüffer \(\ddagger\)†

Thin oxide films with high dielectric constants, often referred to as high-\(k\) dielectrics, are of foremost importance for the complementary metal oxide semiconductor (CMOS) device technology. They have replaced the archetypal gate dielectric material SiO\(_2\) in the field effect transistor (FET) – the heart of central processor units (CPU), in the metal–insulator–metal (MIM) capacitors of the volatile dynamic random access memory (DRAM), and in the floating gate FET of the non-volatile flash memory devices.\(^6\) This step ensured the continuation of the exponential increase of transistor packing density in integrated circuits beyond the 45 nm technology, lowering the power consumption and increasing the operation speed in every subsequent generation of devices.\(^7\)

The substitution of SiO\(_2\) became necessary because at a layer thickness below ca. 1.2 nm, the quantum tunneling effect resulted in an unacceptably high leakage current, and consequently, higher power losses and poor device performance. Materials with higher dielectric constants compared to SiO\(_2\) enabled reduction of the equivalent oxide thickness (EOT) to the subnanometer range at higher physical thickness, allowing to further reduce the transistor size. For instance, in the 14 nm technology an EOT of 0.9 nm was achieved by a high-\(k\) dielectric layer thickness of 2.6 nm, whereas in the upcoming 3 nm technology the targeted EOT of 0.6 nm requires a dielectric layer with a physical thickness below 2 nm.\(^8\)\(^9\) Despite the fact that currently HfO\(_2\), ZrO\(_2\), and Al\(_2\)O\(_3\) are widely used as high-\(k\) dielectrics for CPUs, DRAMs and flash memories, respectively, there is a continuous search for new outperforming materials for these applications.\(^10\)

Rare earth sesquioxides (RE\(_2\)O\(_3\)) have been extensively investigated for applications as high-\(k\) dielectrics.\(^11\)–\(^13\) The favorable combination of high dielectric constants, relatively large bandgaps, high breakdown electric fields, thermal and chemical stability, and small lattice mismatch with silicon promoted this class of materials as attractive candidates for high-\(k\) dielectrics in the CMOS device technology. Among the rare earth sesquioxides, Eu\(_2\)O\(_3\), which is the subject of the present study, demonstrated superior performance as a charge-trapping high-\(k\) dielectric with a layer thickness of 3 nm in low temperature polycrystalline silicon thin-film transistor (LTPS-TFT) nonvolatile memory devices.\(^14\)

The spatial confinement of atoms in nano- and subnanometer-thick layers profoundly alters their vibrational dynamics.\(^15\) Via phonon–phonon,\(^16\) electron–phonon\(^17\) and spin–phonon\(^18\) interactions, these size effects impact important
properties of nanomaterials like thermal conductivity, superconductivity, charge mobility, spin polarization and others, manifesting themselves in phenomena such as self-heating,\textsuperscript{19} increase of the critical temperature of superconductors\textsuperscript{20–22} and spin relaxation,\textsuperscript{23} which play a crucial role in micro-, optoelectronic and spintronic device technologies.\textsuperscript{24–26}

Common features of the phonon density of states (PDOS) of thin films are enhancement of the number of states at low and high energy as well as broadening and shift of the peaks.\textsuperscript{27} Extensive studies revealed that the origin of these vibrational anomalies is tightly connected to the crystal structure and thickness of the layer. In epitaxial thin films and superlattices, the broken translational symmetry at the surface and interface constitutes a source of novel vibration dynamics, which relaxes towards the bulk behavior within a few atomic layers.\textsuperscript{28–38} In amorphous thin films, the structural disorder induces a reduction and a broad distribution of the mean force constants, leading to a significant enhancement of the number of low- and high-energy states.\textsuperscript{39–41}

Fundamentally, the origin of the high-$k$ property is directly related to the vibrational spectrum of a solid, since the static dielectric constant is a sum of the electronic and lattice contributions. While the former equals to the square of the refractive index and thus is intrinsically limited to low values, the high-$k$ values originate from the lattice component, which is inversely proportional to the square of the frequency of transverse optical phonons.\textsuperscript{4,42} This fact, combined with the observed modifications of the PDOS of ultrathin films, calls for a study of the lattice dynamics of high-$k$ dielectrics with thicknesses of several atomic layers.

In this Communication we present a systematic lattice dynamics study of ultrathin Eu$_2$O$_3$ films with thicknesses of 2.2, 1.3 and 0.8 nm deposited on YSZ(001) substrates. The Eu-partial PDOS, derived from nuclear inelastic scattering (NIS) on the M"ossbauer-active isotope $^{151}$Eu, unveils strong size effects that significantly alter the vibrational thermodynamic and elastic properties of the thin films in comparison to a 21.3 nm thick film. Our analysis reveals that these effects primarily originate from an ultrathin EuO layer formed between the thin Eu$_2$O$_3$ film and the YSZ(001) substrate. In addition to the spatial confinement and structural disorder, we demonstrate that the formation of sub-nanometer-thick layers with exotic crystal phases at the interface constitutes another source of vibrational anomalies of thin films and multilayers.

Eu$_2$O$_3$ films with thicknesses of 21.3, 2.2, 1.3 and 0.8 nm, hereinafter referred to as S1, S2, S3 and S4, were deposited on YSZ(001) substrates in the UHV system,\textsuperscript{43} located at the Nuclear Resonance Beamline ID18 (ref. 44) of the ESRF in Grenoble, France, with a base pressure of 5 $\times$ 10$^{-10}$ mbar. After deposition of the Eu metal, the sample was annealed at the same conditions for 60 min in order to achieve a complete oxidation of the Eu metal and to avoid the formation of other oxide phases. To protect the sensitive Eu$_2$O$_3$ films and to mimic the spatially confined geometry of the gate dielectric in a planar FET, the samples were covered at 298 K by 4.0 nm thick Nb layer.

The $^{151}$Eu-partial PDOS of the samples was obtained\textsuperscript{45} from the energy dependence of the probability for nuclear inelastic absorption\textsuperscript{46–48} of X-rays with an energy of 21.5 keV with an energy resolution of 1.1 meV (fwhm).\textsuperscript{48} All samples were illuminated at a grazing angle of 0.15 degree using a focused X-ray beam with dimensions: vertical $\times$ horizontal $= 10 \times 100$ $\mu$m$^2$.

The Eu-partial PDOS of the investigated samples are presented in Fig. 1(a). The PDOS of S1 exhibits a broad peak at around 13 meV, a peak at 24 meV and a high-energy cutoff at

Fig. 1 (a) Eu-partial PDOS of the investigated samples. The solid gray line stands for the ab initio calculated PDOS of a bulk Eu$_2$O$_3$ crystal with cubic symmetry.\textsuperscript{49} The dashed line corresponds to the gray-line PDOS, convoluted with the DHO function with a quality factor $Q = 5$ (see Table 1). The dotted line stands for the Eu-partial PDOS of an EuO film with a thickness of 2.0 nm.\textsuperscript{37} The solid red lines depict the model function described in the text. (b) Reduced PDOS (PDOS/$E^2$) of S1–S4 and of Eu$_2$O$_3$ crystal with cubic symmetry (gray line).
27.5 meV. The gray solid line depicts the ab initio calculated Eu-partial PDOS of an Eu$_2$O$_3$ crystal with cubic symmetry. A comparison with the theory reveals a good agreement with the number and positions of the phonon states, as well as an excess of states at energies up to 12 meV in the experimental PDOS. A reduction of the film thickness to 2.2 nm in S2 leads to an increase of the number of low-energy states and a significant broadening of the spectral features. The effect is particularly strong for the high-energy peak, which practically diminishes. The tendency of enhancement of the number of low-energy states is clearly present in the PDOS of S3 (film thickness of 1.3 nm) and S4 (film thickness of 0.8 nm).

The differences between the samples in the low-energy range of the PDOS is illustrated in Fig. 1(b), where the reduced PDOS (PDOS/E$^2$) is plotted. This figure shows that the number of states between 5 and 7 meV in S2 and S3 exceeds that of S1 in average by factors of 1.6 and 2, respectively, whereas in S4 the excess reaches a factor of 2.3. Compared to the theoretical Debye level of an Eu$_2$O$_3$ crystal with cubic symmetry (gray line in Fig. 1(b)), the number of low-energy states in S1 is increased by a factor of 1.8.

The increase of the number of low-energy states is a generic feature of the PDOS of thin films. At epitaxial surfaces and coherent interfaces, this feature originates from surface- and interface-specific vibrational modes arising from the reduced mean force constants which is a consequence of the broken translational symmetry. In amor- phous thin films, the broad low-energy peak originates from the reduced mean force constants with a large width of distribution associated with the structural disorder. In addition to these well studied effects, below we unveil another potential source of anomalies in the lattice dynamics of thin films and multilayers.

The crystal quality, thickness and composition of the films were determined by X-ray diffraction (XRD) and X-ray reflectivity (XRR) measurements using Cu-K$_\alpha$ energy, and by X-ray absorption near-edge structure (XANES) spectroscopy on the L$_1$ edge of Eu (6977 eV) at the SUL-X beamline of the KIT Light Source. The FitSuite, ATHENA and ARTEMIS codes from the IFEFFIT package were used for reduction and modeling of the experimental data. Fig. 2(a) shows the XANES data obtained from samples S1–S4 along with a reference 100 nm thick EuO film and a commercial Eu$_2$O$_3$ powder sample measured in the same experiment. The XANES data show a distinct difference in the position of the absorption L$_1$ edge of Eu in Eu$_2$O$_3$, where the Eu atoms are in an oxidation state +3, and in EuO where the atoms, similarly to metallic europium, exhibit an oxidation state +2. Unexpectedly, Fig. 2(a) reveals, in addition to Eu$^{3+}$, the presence of Eu$^{2+}$ in the investigated samples. The relative amount of this phase increases with decreasing the film thickness. The presence of metallic Eu can be excluded due to the high partial oxygen pressure in the growth chamber, the elevated temperature of the substrate and the prolonged annealing time. Therefore, the Eu$^{2+}$ peak indicates the formation of EuO, which however, is unexpected since the optimal values for the epitaxial growth of EuO on YSZ(001) were exceeded by 150 K for the substrate temperature and by one order of magnitude for the partial oxygen pressure. The relative Eu$^{2+}$ and Eu$^{3+}$ fractions in the samples were obtained by modeling the experimental data with a linear combination of the XANES spectra of the reference samples. The best-fit results, plotted with solid/red lines in Fig. 2(a), show that this model satisfactorily reproduces the experimental data. The obtained relative amount of the EuO phase and the corresponding film thickness (assuming a homogeneous layer) are summarized in Table 1.

The formation of an EuO layer at the interface with the YSZ(001) substrate is further confirmed by the analysis of the XRR data presented in Fig. 2(b). The red lines correspond to the best-fit results assuming an EuO layer formed between the Eu$_2$O$_3$ film and the YSZ(001) substrate (see inset Fig. 2(b)). The total thickness of the oxide layer (Eu$_2$O$_3$ + EuO), the thickness and the corresponding relative volume fraction of the EuO interface layer are presented in Table 1. While the results from

Fig. 2 XANES (a) and XRR (b) data of the investigated samples. The red lines mark the corresponding simulations (see text for details). The inset in (b) depicts the model used for the fit of the XRR data.
both methods are in a qualitative agreement on the increasing contribution of the EuO phase as the film thickness is reduced, the obtained relative contribution of this phase from the XANES experiment systemically exceeds that derived by the XRR measurement. Yet the thickness of this interface layer obtained by both methods is in the sub-nanometer range except in S1 where it reaches 1 nm.

A combined experimental and theoretical study revealed strong confinement effects in the lattice dynamics of thin EuO films. Namely, the Eu-partial PDOS of an EuO film with a thickness of 2.0 nm exhibits a broad peak at 10.5 meV and a high-energy cutoff at 25 meV. These features are very similar to those exhibited by the PDOS of S2–S4 (Fig. 1(a)). In order to thoroughly explain the experimental observations, the data were modeled by a superposition of the ab initio calculated Eu-partial PDOS of Eu2O3 with cubic symmetry \( g_\text{Eu}(E) \) (gray line in Fig. 1(a)) and the experimentally determined PDOS of a 2.0 nm thick EuO film, \( g_\text{Eu}(E) \) (dotted line in Fig. 1(a)), representing the interface. Furthermore, the broadening of PDOS features is a common characteristic for nanoscale materials that originates from phonon scattering at atoms located at irregular lattice sites, i.e., grain boundaries, defects and dislocations at surfaces and interfaces, as well as within the nanostructure. This effect is satisfactorily described by the damped harmonic oscillator (DHO) function, which introduces an energy-dependent broadening of the phonon features reflecting the fact that high-frequency phonons scatter more efficiently. The strength of phonon damping is inversely proportional to the quality factor \( Q \) of the DHO function. To account for these two sources of anomalies in the PDOS of the thin films, namely the phonon damping and the presence of an EuO interface layer, the following model function is defined:

\[
g_\text{model}(E, Q) = A g_\text{Eu}(E) + (1 - A) g_\text{th}(E, Q)
\]

with \( A \) being the relative EuO interface atomic fraction. The experimental PDOS of S1–S4 were modeled with the function \( g_\text{model}(E, Q) \) using the least-squares method with \( A \) and \( Q \) being variable parameters. The results are plotted with red/solid lines in Fig. 1(a) and show that the used model satisfactorily reproduces the experimental data. The obtained values of \( A \) and \( Q \) are summarized in Table 1. The relative interface fraction falls in between the values obtained by XRR and XANES, which indicates the consistency of the results obtained by three independent experimental techniques. 

| Sample | Thickness (nm) | XRR (%/nm) | XANES (%/nm) | NIS (%/nm) | \( Q \) |
|--------|---------------|------------|--------------|------------|------|
| S1     | 21.3          | 4/0.9      | 5/1.1        | 6/1.3      | 12   |
| S2     | 2.2           | 18/0.4     | 35/0.77      | 23/0.51    | 4    |
| S3     | 1.3           | 38/0.5     | 47/0.61      | 45/0.59    | 4    |
| S4     | 0.8           | 37/0.3     | 56/0.45      | 51/0.41    | 5    |

Thickness of the oxide layer (Eu2O3 + EuO) and relative contribution (A)/thickness of the EuO interface layer obtained by the corresponding methods. \( Q \) is the quality factor of the DHO function (see text for details).

The observed broadening of the Eu-partial PDOS of the ultrathin films, quantified with the quality factor \( Q \), most likely applies as well for the vibrational modes of the oxygen atoms, characterized with higher frequencies and being primarily responsible for the high-\( k \) property of the REO3. This suppression and even disappearance of transverse optical modes could potentially reduce the lattice contribution to the dielectric constant. Indeed, a decrease of the static dielectric constant of La2Zr2O7 thin films compared to the bulk material was observed and explained by a broadening and vanishing of the infrared-active phonon modes due to structural disorder of the thin films.

The reduction of the film thickness has important implications on the elastic and thermodynamic properties. Table 2 summarizes the experimental and theoretical values of the mean force constant \( F \), mean square atomic displacement \( \langle x^2 \rangle \), vibrational entropy \( S_v \) and lattice specific heat at constant volume \( C_V \) obtained from the Eu-partial PDOS of an EuO crystal with cubic symmetry (calculated for the experimental lattice constant of 10.80 Å) for samples S1–S4 and of a 2.0 nm thick EuO film. A decrease of \( F \) in S2 by 6.5% compared to S1 results in an increase of \( \langle x^2 \rangle \) by 17.6%. A further softening of the EuO layer thickness is also consistent with the values derived by the other two methods. The values of \( Q \) are abruptly reduced from 12 in S1, corresponding to an almost bulk-like crystal (compare the width of the peak at 25 meV in the experimental and theoretical PDOS) to 4 in S2 and S3 and 5 in S4 (this difference we attribute to the statistical error). These values of \( Q \) unveil strong and very similar phonon damping in the thin Eu2O3 films, despite the fact that the total oxide layer thickness varies by a factor of 2.8 from S2 to S4. This result indicates that the investigated Eu2O3 ultrathin films exhibit crystal lattices of similar quality (as confirmed by the XRD study). 

A closer comparison of the experimental PDOS to the \( g_\text{model}(E, Q) \) (Fig. 1(a)) demonstrates that the model function systematically underestimates the number of phonon states between 3 and 10 meV. Most likely this difference arises from vibrational states of Eu atoms located at the Nb/Eu2O3 interface, which was not included in the model.

| Sample | \( F \) (N m\(^{-1}\)) | \( \langle x^2 \rangle \) (Å\(^2\)) | \( S_v \) (k\(_B\) per atom) | \( C_V \) (k\(_B\) per atom) |
|--------|----------------------|-------------------|-----------------|-----------------|
| Eu2O3 bulk\(^{49}\) | 208 | 0.01237 | 4.462 | 2.870 |
| S1     | 168(5) | 0.017(1) | 4.68(2) | 2.89(2) |
| S2     | 157(4) | 0.020(1) | 4.86(2) | 2.90(2) |
| S3     | 133(4) | 0.023(1) | 5.09(2) | 2.92(2) |
| S4     | 123(3) | 0.024(1) | 5.19(2) | 2.93(2) |
| EuO film\(^{37}\) | 89(7) | 0.031(2) | 5.58(4) | 2.94(3) |
lattice by 20.8% in S3 and by 26.8% in S4 increases ($\chi^2$) by 35.3% and 41.2%, respectively. $S_1$ increases by 3.9% in S2, 8.8% in S3 and 10.9% in S4 compared to S1, whereas $C_6$ shows only a marginal enhancement by 1.4% in S4. Since the quality factor of the DHO function remains essentially the same in S2–S4 (Table 1) and the EuO film exhibits significantly different properties from those of bulk Eu$_2$O$_3$ (Table 2), the observed anomalous behavior of the thermoelastic properties of the films originates primarily from the increase of the EuO interface fraction as the film thickness is reduced.

In summary, using nuclear inelastic scattering of synchrotron radiation on the Mössbauer-active isotope $^{151}$Eu, we determined the Eu-partial phonon density of states in Eu$_2$O$_3$ films with thicknesses of 21.3, 2.2, 1.3 and 0.8 nm deposited on YSZ(001). The PDOSs exhibit broadening and suppression of the peaks accompanied by up to a four-fold enhancement of the number of low-energy states in the thinnest layer compared to the ab initio calculated PDOS of a perfect Eu$_2$O$_3$ crystal. Our analysis demonstrates that the former effect, which could potentially reduce the dielectric constant of ultrathin high-k films, arises from the presence of lattice defects and grain boundaries and reflects the reduced phonon lifetimes observed in thin films. The structural characterization unveiled the formation of a sub-nanometer-thick EuO layer at the interface between the Eu$_2$O$_3$ film and the YSZ(001) substrate. The spatially confined atomic vibrations of this interface layer contribute significantly to the increased number of low-energy phonon states and to the anomalies in the thermodynamic and elastic properties of the investigated samples.

The presented results demonstrate that in addition to spatial confinement, anisotropy, surface- and interface-specific vibrational modes, sub-nanometer-thick layers exhibiting exotic crystal phases that might form at the interface between two materials constitute a source of novel vibrational dynamics of thin films and multilayers. Moreover, the targeted embedment of such ultrathin layers might be a successful approach towards ultimately engineered heterostructures for phononic applications.

Conflicts of interest

There are no conflicts of interest to declare.

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