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S.I.: COMPUTATIONAL ELECTRONICS OF EMERGING MEMORY ELEMENTS

**A computational study of hafnia-based ferroelectric memories: from ab initio via physical modeling to circuit models of**   
**ferroelectric device**

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**Abstract** The discovery of ferroelectric properties of binary oxidesrevitalizedtheinterestinferroelectricsandbridgedthe scaling gap between the state-of-the-art semiconductor tech-nology and ferroelectric memories. However, before hitting the markets, the origin of ferroelectricity and in-depth studies of device characteristics are needed. Establishing a correla-tion between the performance of the device and underlying

presented and potential further applications of ferroelectric devices are outlined.

**Keywords** Modeling · FRAM · FeFET · Wake-up ·Ferroelectric HfO2 · Ferroelectric memory

physical mechanisms is the first step toward understanding **1 Introduction**

the device and engineering guidelines for a novel, supe-

rior device. Therefore, in this paper a holistic modeling approaches which lead to a better understanding of ferro-electric memories based on hafnium and zirconium oxide is addressed. Starting from describing the stabilization of the ferroelectricphasewithinthebinaryoxidesviaphysicalmod-eling the physical mechanisms of the ferroelectric devices are reviewed. Besides, limitations and modeling of the multilevel operation and switching kinetics of ultimately scaled devices as well as the necessity for Landau–Khalatnikov approach are discussed. Furthermore, a device-level model of ferro-electric memory devices that can be used to study the array implementation and their operational schemes are addressed. Finally, a circuit model of the ferroelectric memory device is

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In the age of mobile devices nonvolatile memories have become an essential part of every electronic system. Tra-ditionally, nonvolatile memories are limited with respect to rewrite speed and cycling endurance due to the so-called voltage time dilemma [1]. Additionally, the high voltages required during the rewrite process make their integra-tion into scaled complementary metal-oxide-semiconductor (CMOS) processes more and more complicated. The stable remanent polarization at zero electric field together with the fact that the polarization can be switched by an electrical field makes ferroelectrics a natural choice to realize a nonvolatile memory [2,3]. Therefore, already in the 1950s the concept of a ferroelectric memory was proposed [4] and first attempts were made to realize solid state memories based on the ferro-electric Bariumtitanate [5,6]. However, these attempts were based on pure cross-point arrays and the resulting disturb issues could not be solved. The rapid development of semi-conductortechnologyenabledtosolvethisfundamentalissue

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| 1 | NaMLab gGmbH, Noethnitzer Str. 64, 01187 Dresden, | byaddingaMOSselecttransistortothecellarchitecture.Asa |
| 2 | Germany | result the first commercial ferroelectric memories appeared |
| Department of Applied Sciences and Mechatronics, Munich | in the early 1990s [7]. This success spurred the hope that |
| 3 | University of Applied Sciences, Munich, Germany | within a few years a high density memory that would have |
| Ferroelectric Memory GmbH, 01187 Dresden, Germany | the cell size and performance of DRAM and at the same time |
| 4 | Chair of Nanoelectronic Materials, Institute of | be nonvolatile would emerge [8]. Soon, however, it was rec- |
| ognized that the integration of very complicated perovskite |
| Semiconductors and Microsystems, TU Dresden, Dresden, |
| Germany | or even layered perovskite materials requires a large integra- |

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tion effort that would hinder a scaling on the same pace as CMOS and related memory concepts based on floating gates or charge trapping layers [9]. Since the readout of a ferroelec-tric capacitor is based on the charge that is transferred during switching [1] a 3-dimensional capacitor would be required below 100 nm groundrules. This is adding new integration challenges for the complex materials not only of the ferro-electric itself but also the required metal-oxide electrodes. For this challenge no solution can be provided yet [10,11]. Therefore, the development of such perovskite-based mem-ories were successful but at the same time seem to have hit a wall at about the 100 nm groundrules [12]. Another path to scale the memory cell without going into the third dimen-sion is the integration of the functional ferroelectric film into a field-effect transistor. Such an approach, called the ferro-

**2 Engineering the material: ab-initio study**

The observation of ferroelectricity in thin hafnia films has been surprising since the most stable monoclinic P21/c phase, the high-temperature tetragonal P42/nmc and cubic Fm3m phases and the high pressure orthorhombic Pbca phase (the well-known equilibrium phases) are centrosymmetric and nonpolar. Following older experimental [24] and theoreti-cal [25] work, the metastable, polar, orthorhombic Pca21 was proposed as the ferroelectric phase, which is nearly indistinguishable from tetragonal in grazing incidence X-ray diffraction (GIXRD). With a combined transmission electron microscopy (TEM) and nanoscale electron diffraction study, Sang et al. [26] were able to identify the Pca21 phase in a thin, ferroelectric hafnia film with some uncertainty with regard

electric field-effect transistor (FeFET) was proposed already to the Pbca phase.

in the late 1950s [13,14]. However, perovskite or layered perovskite ferroelectrics like lead–zirconate–titanate (PZT) and strontium–bismuth–tantalate (SBT) in contact with sil-icon require an interface buffer layer to avoid silicidation reactions. Due to the high permittivity of perovskites it is nearly impossible to match the permittivity of the ferroelec-tric with a nonferroelectric interface layer while trying to reduce the gate stack thickness. An internal depolarization field under zero applied voltage is the consequence leading to poor data retention [15]. Using hafnium oxide as the inter-face layer between a very thick SBT layer and silicon [16], nonvolatile retention could be demonstrated. However, the thickness requirements for the SBT layer hinder the scaling of such devices to the deep sub-100 nm regime as well.

The discovery of ferroelectricity in hafnium oxide by Boescke et al. [17] may resolve these historic issues of ferro-electric memory devices explained above. Hafnium oxide

Density functional theory (DFT) total energy calculations qualitatively mostly agree that the Pca21 phase of hafnia is the second most stable phase with about 60 meV/f.u., fol-lowed by the tetragonal phase with 95 meV/f.u. (relative to monoclinic). The considerably different values for the energy differences result from different density functionals where the local density approximation (LDA) typically gives smaller and the generalized gradient approximation (GGA) larger energy differences. Further theoretical uncertainty arises from d-orbitals in standard DFT which can be better described with more sophisticated methods like DFT+U or HSE (some examples are shown in Fig. 1a). Results obtained with this method are consistent and lead to conclusion that the ferroelectric phase has to be stabilized relative to the mon-oclinic phase by certain mechanisms. Important for phase stability is the expression of the free energy

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| is the standard gate dielectric in sub-45 nm CMOS pro- | *F* = *U* − *T S,* | (1) |
| cesses. Proven fabrication and integration schemes exist, and |

hafnium oxide is thermodynamically stable on silicon [18]. Therefore, this discovery has drastically changed the view on ferroelectric memories [19]. Both the scaling of classi-cal capacitor-based memories as well as the realization of FeFET-basedmemoriesseemtobeinreachagain[20,21]and even the possibility of making classical DRAM nonvolatile has been pointed out recently [22,23]. However, in order to apply the unexpected ferroelectric behavior of hafnium oxide to reliable products a number of fundamental questions need to be answered: the origin of the ferroelectric phase, the con-trol of the influencing process parameters, the device and array concepts as well as the degradation under use condi-tions. With such fundamental questions to be addressed, the

where *U*0 in *U* = *U*0+*U*zero is the total energy, *T* the temper-ature,*U*zero is from zero point vibrations and S the vibrational entropy. Dependence on electric field, stress and surface energy will be discussed in Sect. 2.1. Figure 1a shows that the vibrational entropy lowers especially the free energy of the tetragonal phase with temperature, which should therefore prevail under annealing conditions. For zirconia, the energy difference between the competing ferroelectric and tetrago-nal phase becomes very small and the free energy (Fig. 1b) nearly vanishes. In the solid solution Hf1−*x*ZrxO2 the free energy differences can be linearly interpolated between HfO2 and ZrO2. Thus, ZrO2 can be considered as a dopant sta-bilizing the ferroelectric phase, although with a competing

modeling on all abstraction levels from atomistic material tetragonal phase.

simulations via device simulations up to the array and circuit level are required. This paper summarizes the recent progress on this exciting topic.

In summary, this result describing the ferroelectric phase as a metastable phase matches well with the fact that it only exists under certain circumstances as there are film thickness and grain size, dopants, impurities and as speculated stress

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| 200 **(a)**  **(b)**  further stabilization mechanism in addition to the dopant is   |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | | |  |  | | --- | --- | | erences (meV/f.u.) | 150  100 | |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | |  |  | | --- | --- | | /f.u. | 80 | |  |  |  |  |  |  |  |  |  |  |  |  |  | |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | V |  |  |  |  |  |  |  |  |  |  |  |  |  | |  |  |  | T | ot | a |  |  |  |  |  |  |  |  |  |  |  | e |  |  |  |  |  |  |  |  |  |  |  |  |  | |  |  |  | T | o | a |  |  |  |  |  |  |  |  |  |  |  | m |  |  |  |  |  |  |  |  |  |  |  |  |  | |  |  |  | F | e | e |  |  |  |  |  |  |  |  |  |  |  | ( |  |  |  |  |  |  |  |  |  |  |  |  |  | |  |  |  | F | e | e |  |  |  |  |  |  |  |  |  |  |  | |  |  | | --- | --- | | s | 60 | |  |  |  |  |  |  |  |  |  |  |  |  |  | |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | rence |  |  |  |  |  |  |  |  |  |  |  |  |  | |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | e |  |  |  |  |  |  |  |  |  |  |  |  |  | |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | ff |  |  |  |  |  |  |  |  |  |  |  |  |  | |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | di |  |  |  |  |  |  |  |  |  |  |  |  |  | | |  |  | | --- | --- | | Energy diff | 50 |   0 |  |  |  | |  |  |  |  |  | |  |  |  | |  |  | |  |  | | --- | --- | | ree energy | 20  0 | | |  |  |  |  |  |  | |  |  |  | | | | |  |  |  |  |  | |  |  |  |  |  | |  |  |  |  |  |  |  |  |  | | f | | t | | | f | | t | | f | | t | | f | | t | |  | f | | t | f | t | | f | | t | f | t | |  | | LDA [A] | | | | | LDA [B] | | | | GGA | | | | HSE | | | |  | - | | | 6.25%Vo | | | - | | | 6.25%Vo | | |  | |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | HfO2 | | | | | | ZrO2 | | | | | |  |   was already proposed in [33] to explain the stabilization of  the tetragonal phase in thin films by DFT data correctly.  required to avoid excessive leakage currents and charge  Charge compensation in doped, ferroelectric hafnia  involved. We believe that surface or interface energy in poly- crystals, discussed in Sect. 2.2, is a major effect closing the  gap between simulation and experiment. Such a mechanism | |

**Fig. 1 a** Total and free energy differences of HfO2 ferroelectric (*f* ) and tetragonal (*t*) phase relative to monoclinic for different DFT methods, **b** free energy of ZrO2 and HfO2 as well as the effect of 6% oxygen vacancies *(V*O*)* on free energy. LDA [A] values are from Ref. [30], LDA [B] from [29], GGA from [31] and HSE from [32]. Values for (**b**) are calculated based on Ref. [29]

and or electric fields. In pure hafnia, the ferroelectric phase free energy is about 50 meV/f.u. relative to the monoclinic ground state such that it can be only stabilized in extremely thin films [27]. On the other hand, in pure zirconia the tetrag-onal phase is only a few meV/f.u. close to the ferroelectric phase such that an external electric field is sufficient to induce its stability [28,29]. Around the 50% mixture of hafnia and zirconia the ferroelectric phase is very stable under thin film conditions (up to a thickness of about 25 nm).

Hf1−*x*Zr*x*O2 is a model system for ferroelectric phase sta-bilization with a dopant. Namely, for a thin film (of about 10nm),increaseintheconcentrationofanappropriatedopant for a few anionic % lowers the free energy of the ferroelectric phase as well as of the tetragonal and cubic phases relative to the monoclinic one. For the four valent Si there is a con-centration range from 3–5 cat% (cationic ratio Si/(Hf+Si)) favoring the stabilization of the ferroelectric phase. For the

build-up, might work in pairing dopants with oxygen vacan-cies, but still needs to be demonstrated in detail. By calculat-ing the occupation states of the defect levels in the band gap, the efficacy was shown for monoclinic hafnia and Y [35], Al [36], Gd [37] or Ba [38] as a dopant.

Oxygen vacancies, *V*O, in hafnia without compensating dopants are causing leakage current and charge build-up. The defect structure of the monoclinic [39] phase has been researched intensively and reveals the double-charged *V*++ O with unoccupied states about 1eV below the conduction band as the most stable structure for zero applied field.

Not only the paired, but also the unpaired vacancies might contribute to the desired phase stabilization. Here, some pre-liminary results have been published [40], shown in Fig. 1b. 6.25 ani.-% *V*O in HfO2 lowers the ferroelectric phase by 15 meV/f.u. and the tetragonal phase by 20 meV/f.u. rela-tive to monoclinic phase. Interestingly, introduction of *V*O favors the stabilization of the tetragonal phase with respect to the ferroelectric. This leads, leading to the typical concen-tration window with the tetragonal phase dominating at large concentrations. If oxygen vacancies play a role in the phase transformation observed during field cycling, however, they have to be present somewhere with a concentration of a few percent (not all *V*O-s are active from the electrical point of

three valent dopants N, Al, Y, Gd and La and for the two view).

valent Sr and Ba [32], concentration ranges from few to more

than 10 cat% have been found. Beyond this range, the film becomes tetragonal for the four valent and cubic for the three and two valent dopants. Even though this topic is well suited for DFT simulations, apart from Fischer et al. [33] and Lee et al. [34] (where dopants stabilizing the tetragonal high-*k* phase were searched), no systematic study of phase stabiliza-tion with dopant has been reported so far. There are several reasons for the slow progress, whereas tetragonal or cubic phase stabilization clearly dominates beyond some doping range, ferroelectric phase stabilization seems to be possible only in a distinct concentration window requiring more accu-rate calculations. The introduction of various dopants might be accompanied by the formation of oxygen vacancies pro-viding the charge compensation, and consequently requiring the calculation of a large number of structures to find the most favorable configuration. Finally, the resulting total energy or more demanding free energy calculation results do not fully explain the phase stabilization in contrast to the experiments. Obviously, either the DFT calculations are wrong, or some

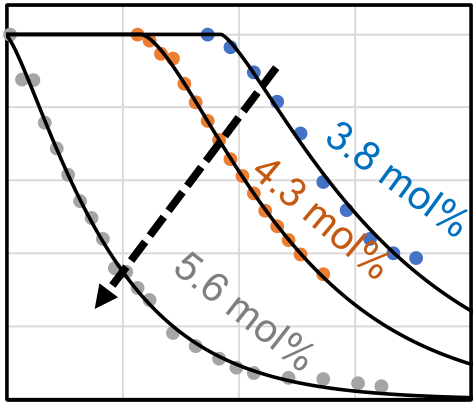
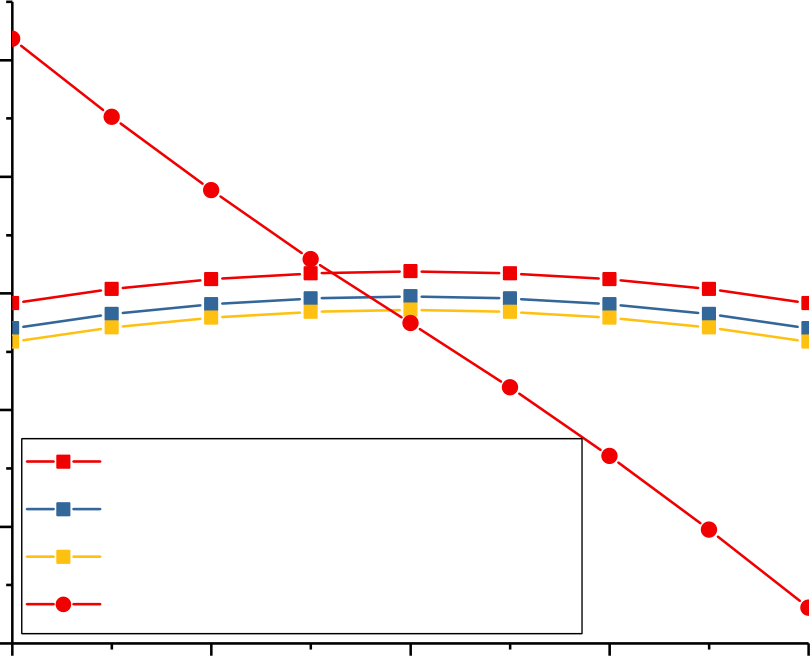
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**2.1 Phase transitions within binary oxides**

A phase transition of ferroelectric hafnia into the tetragonal phase is desired when pyroelectric or electrocaloric effects are being exploited, controlled by temperature *T* and electric field *E* in a thermoelectric cycle. In contrast to the pyro-electric and electrocaloric applications, phase transitions are highly undesired in memory application. Nonetheless, for reaching the desired phase stability it is important to investi-gate its dependencies. The free energy for each phase *φ* can be calculated as

*Fφ* = *Uφ* − *T Sφ* + *�*0*,φ Dφ Eφ* + *�*0*,φσi j,φs ji,φ* + *�φ,* (2)

where *�*0*,φ* denotes the volume of a formula unit of the phase, *Eφ* the electric field, *D* the dielectric displacement, S the vibrational entropy, *σ*ij and *s*ij stress and strain tensor, respectively, and *�φ* a surface energy contribution.



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| 80 | | | | 2 | 4 | normalized Pr (Pr/Pr\_max) | 1 | Silicon content   increase | | | |
| Free energy [meV/f.u.] | 60 | P42/nmc T=80K | | 0.8 |
| 0.6 |
| 40 | 0.4 |
| 20 | 0.2 |
| 0  100 | | 200 | 300 | 400 | 500 |
| Temperature [K] | | | | | |
| 0 | | P42/nmc T=230K  P42/nmc T=300K | |
| Measured  Simulated | | | | | |
| -20 | | Pbca1 T=80K, T=230K, T=300K | |
| **Fig. 3** Remanent polarization in Si: HfO2 for different doping cat% as | | | | | |
| -4 | | -2 | 0 |
| a function of temperature | | | | | |
| Ez [MV/cm] | | | |

**Fig. 2** Free energy as a function of the electric field in the polarization

direction *E*z and the temperature for the tetragonal and ferroelectric phase of ZrO2 related to the monoclinic phase. The calculated values have been taken form [29]

Thecalculatedtemperaturedependencefrom*Uφ,*zero–TS*φ* with the electric field Ez in the polarization direction is shown in Fig. 2. It can be seen that a 150 K decrease in temperature results in about a 4 meV/f.u. increase in the tetragonal energy difference. This allows the ferroelectric phase stabilization at the phase boundary to the tetragonal phase by lowering the temperature, as shown in [41].

The impact of an electric field on the free energy was stud-ied by Materlik et al. [29] to explain the anti-ferroelectric (AFE) behavior in thin ZrO2 as an electric field-induced phase transformation and by Batra et al. [42] to explore phase stabilization by internal fields. For a field strength of 1 MV/cm the value of *�*0DE is 10 meV/f.u., Fig. 2 shows the calculated scenario of AFE behavior as field-induced phase transition which fits to the observation of [41] in thin zirconia. These findings are of high importance for the stabilization of the distinct material into novel AFE nonvolatile memories

effect to stabilize the ferroelectric phase [30]. The required stress depends on the energy difference value to be overcome (see Fig. 1), which fluctuate for different DFT methods. In [30] it was shown that a planar, compressive strain of 1% (which corresponds to a stress of about 3 GPa) is required for stabilization of the ferroelectric phase. Materlik et al. [29] calculated the required stress to be 8 GPa which seems unre-alistic high and therefore, a surface energy mechanism was proposed. Finally, Batra et al. [42] assumed a combination of 2% compressive strain together with an internal electric field of 1.5 MV/cm. Although stress or strain can play a role in stabilization of the ferroelectric phase in hafnia and zirco-nia, it seems that the first-order effect must lay in some other phenomenon.

So far, we have treated phase stability in the thermo-dynamic picture with the free energy. If large barriers are involved in a transition, kinetic effects may become visible. The transition barrier from the tetragonal to the ferroelec-tric phase has been calculated by several authors [29–31] being about 30 meV/f.u. and of first order. The transition bar-

[22,23]. rier from monoclinic to ferroelectric has not been reported,

From Eqs. (1 and 2) one would expect a temperature induced phase transition to the tetragonal or cubic phase at a fixed temperature (the Curie temperature). In Si doped HfO2 [40] a large interval of transition temperatures has been found (see Fig. 3). This has been explained with the surface or interface energy contribution *�φ* which differs for grains of different size in the film. Since the derivative of *P*r with respect to *T* is the pyroelectric coefficient (sum of proper and giant, phase transition induced contribution), the perspective is to tailor the temperature range and magnitude of the pyro-electric coefficient with Si-doping as well as the grain size

but should be much larger around 210 meV/f.u., accord-ing to [43]. This raises the question whether a field-induced transformation from monoclinic to ferroelectric is possible; however, there is no clear evidence so far.

Polarization reversal is a special case of phase transition. Huan et.al. [31] have proposed that the tetragonal phase is at the intermediate state between both polarities of the ferro-electric phase (Fig. 4).

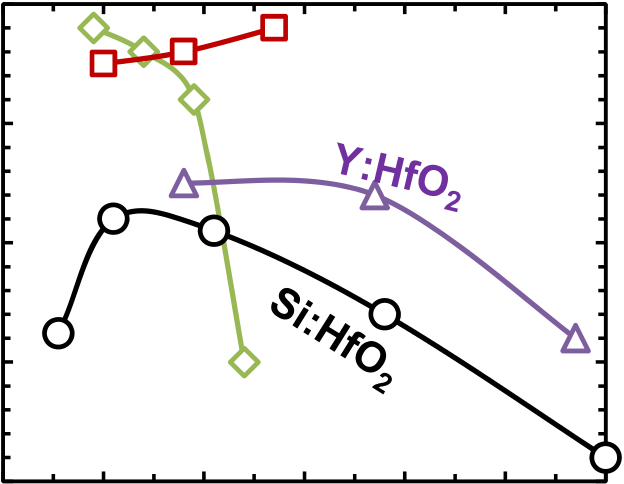
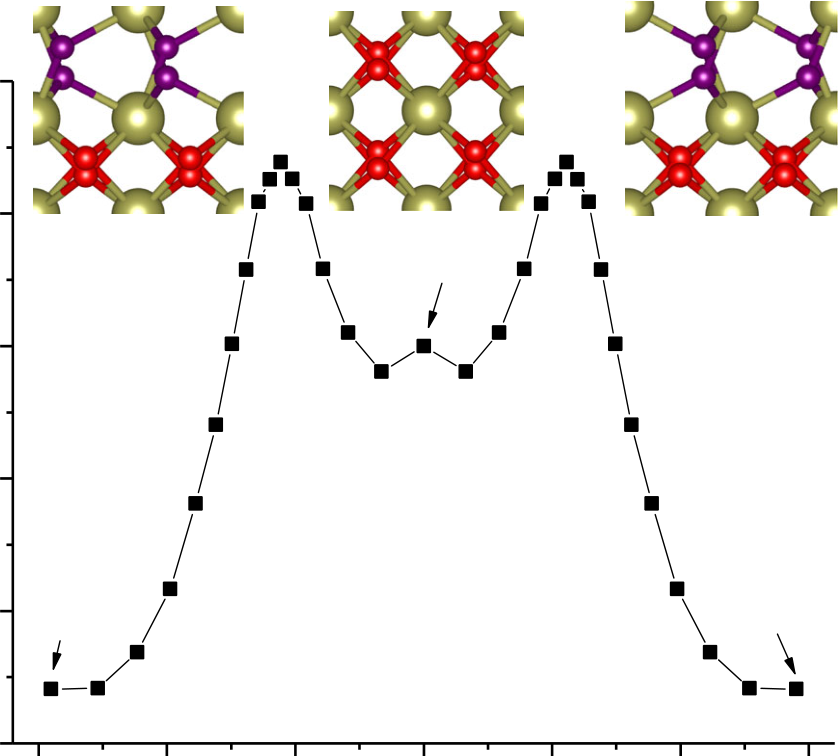
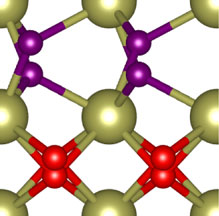
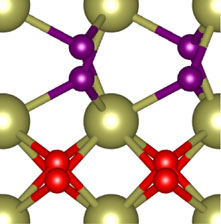
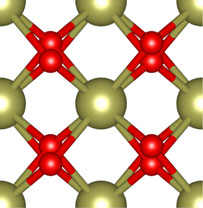
A simple first-order estimate for the coercive field, taking the smaller polarization at the transition state into account [30] results in a value of 1.2 MV/cm which is in good agree-

distribution [40]. ment with the experiments. A more sophisticated approach

A last constituent of Eq. (2) to be discussed is the energy calculating the barrier height under the applied field has not

of stress or strain. Stress has been claimed as the major been reported so far.

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| 40 | | | | | | | | | Pemanent polarization [µC/cm²] | 20 | 0 | 10 | **Gd:HfO2** | | | | 50 | 60 |
| Energy differeneces (meV/f.u.) | 20 | Pca21 | P42/nmc | | | -Pca21 | | | 15 | **HZO** | | | |
| 0 | 10 | 20 | 30 | | 40 |
| 5 |
| -20 |
| 0 |
| -40 | Film thickness [nm]  **Fig. 5** Thickness dependence of *P*r for different doped hafnia  free energy contribution to a grain is | | | | | | | | | |
| -60 | | | | | | | | |
| -3 | | | -2 | -1 | 0 | 1 | 2 | 3 |
| image distance (bohr)  **Fig. 4** DFT calculation of the transition path from the ferroelectric phase via the tetragonal to the negative ferroelectric phase. Energy dif-ferences are produced with pseudopotentials using the String method. The image distance is calculated as the difference of the Frobenius norm of all coordinates | | | | | | | | |
| *�φ* =Surface area | | | *γφ* = | �2 *t*+ 2 | | � | *γφ* | (3) | |
| being a function of *d* as well as *r*. Hence, free energy contri- | | | | | | | | | |

bution to a grain in average is a function of the grain radius   
distribution *f (r, t, x)*, where *x* is the dopant concentration.

All the calculations for kinetic effects reported for hafnia have not considered long range field effects which are present in perovskite ferroelectrics for which effective Hamiltonian models have been developed [44].

**2.2 Stabilization of the ferroelectric phase: surface or**  **interface energy model**

The thickness dependence investigation of the free energy shows that the decrease in the hafnia film thickness favors the ferroelectric phase, whereas the underlying phase in thick films becomes unstable. In contrast to the typical observa-tion in perovskite-based ferroelectrics this trend of binary oxides is very favorable for the integration of ferroelectrics in ultimately scaled devices. It should be noted that there are probably exceptions for the rule that can be concluded from the experimental results shown in Fig. 5. Besides the dopant concentration window of stability there is a maximum thick-ness for which ferroelectric phase in Hf0*.*5Zr0*.*5O2 is stable, with an upper boundary of about 25 nm. Even though there is an indication of decrease in *P*r at film thicknesses of 5.5 nm, however thinner films have not been investigated so far.

A surface or rather interface energy related size effect defining the crystallographic phase of individual grains has been proposed by Materlik et al. [29] and Künneth et al. [45], respectively, assuming cylindrical grains of radius *r*

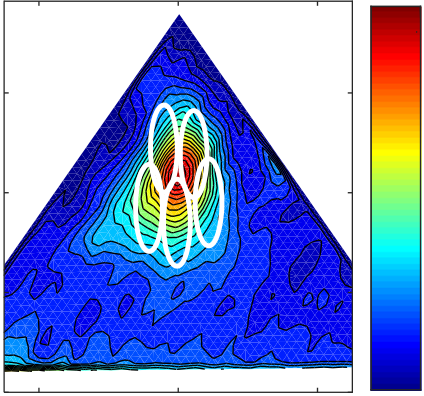
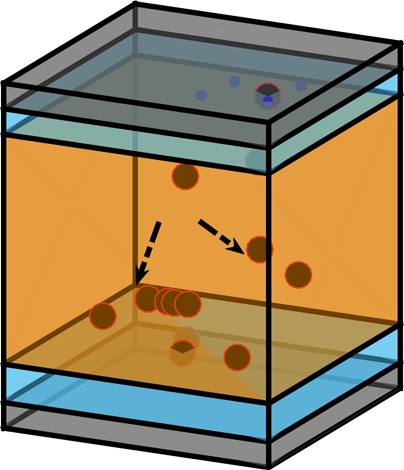
Crucial in the model are the values of surface or inter-face specific energies *γφ*. The suppression of the monoclinic phase with a size effect requires *γ*m *> γ*t and *γ*m *> γ*f (where *γ*m*, γ*t*, γ*f denote the interface or surface specific energy of monoclinic, tetragonal and ferroelectric phase, respectively). The former condition has been experimentally confirmed and the condition *γ*t ≈ *γ*f is reasonable considering the simi-larity of the phases. Furthermore, the term *γφ(x)* depends on the dopant concentration *x*. In [29] a free energy model for Hf1−*x*Zr*x*O2 was developed with a linear dependence of the crystal free energies and *�φ* on *x* (due to the good sol-ubility of HfO2 in ZrO2 and vice versa). Hence, resulting in *x* linear free energies *Fφ(x)* for each phase where the intersections define the phase boundaries. Depending on the hafnia–zirconia mixtures, naturally a window of stable ferro-electric phase in *x* appears, varying with thickness, as shown in Fig. 6. This model is considered to describe the thickness dependence of the ferroelectric phase in Hf1−xZr*x*O2. Gen-erally, for other doped hafnia films, a part of the stabilization is achieved with the dopant and the thickness dependence could be modeled with the surface energy contribution. In [45] the model reported in [29] has been refined with proper inclusion of the grain size distribution. By using total ener-gies from DFT, data can be reproduced quite well, however, the values for *γφ*(x) are taken from a fit. Batra et al. [42] have used DFT calculations for *γφ* (Hf) and obtain a size effect, although no good fit to data, which is no surprise due to the present DFT error present due to the chosen DFT functional

and height *h* = *t* (where *t* denotes the film thickness). The (see Fig. 1 for details).

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| Pr [ | µC/c | m2] |  |
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| **(a)** | | **(b)** | | | **(a)** | **(b)** | | | | |
| **TiN** | **Vacancies** | **HfO** | **2** |  |  | 3 | | | | |
| **IF** |
| **IF** | E c [MV/cm] | 2  1 | | | |
| 0 | | -2 | 0 | 2 |
| **TiN** |
| E | | | bias [MV/cm] | |
| **Fig. 8** 3D geometry of the **a** FeCap structure implemented in MDLab | | | | |

software used for modeling of the vacancy diffusion process and **b** FeCap multidomain model developed in TCAD

techniques. The pristine stage of the lifetime of ferroelectric capacitors is typically characterized by a double current peak feature in the dynamically measured transient current [55, 56]. Since the polarization is the integral of the transient cur-rent, the double current peak characteristics is transformed into a pinched hysteresis loop in the polarization–voltage domain [57]. The continuous stress (field cycling) of the device results in a merging of the double peak characteristics and corresponding opening of the initially, pinched hysteresis loop [56]. During that stage, it was reported that the leakage current stays rather constant [55], whereas the transmission electron microscopy investigations pointed on phase transi-tions within the material [55,58]. Further cycling of the FE capacitor results in a broadening of the current peak and as a consequence decrease in the *P*r amplitude (fatigue stage) and is followed by the increase in the leakage current magnitude

**Fig. 9 a** SEM top-view of a polycrystalline hafnia film: granular struc-ture of the capacitor device where each FeCap (*sketched circled area*) has a random grain distribution. **b** Corresponding domain separation depicted on a Preisach density plot obtained using a FORC technique. Dark red and blue denote maximum and minimum of the Preisach den-sity, respectively (Color figure online)

a tetragonal phase. These two factors significantly impact the voltage distribution inside the dielectric layer. Hence, a superposition of the lower dielectric constant and the elec-tron trapping at interfacial defects can results in either a local increase or decrease in the effective field, which influences the resulting polarization of the ferroelectric film.

In the next step, the polycrystallinity of the HfO2 layer has to be explicitly taken into account. In order to stabi-lize the desired crystal phase and therefore the ferroelectric properties, hafnia-based devices are crystallized using an 800–1000◦C anneal for 20 s in nitrogen atmosphere which induces a granular, polycrystalline morphology of the film.

[55,59]. Scanning electron microscopy micrographs (see Fig. 9a)

In the following a review of the structural and physical properties that have to be accounted for in the model is given. First the impact of the two additional interfacial regions on theelectricalbehavioroftheMIM(TiN/X:HfO2/TiN)capac-itor (where X is the dopant) has to be considered. In addition to the TiO*x*N*y* and TiO*x* interfacial layer formation due to oxygen scavenging from the HfO2 and nitrogen diffusion, results of the TEM study [55,58] revealed parasitic tetrago-nal regions of the HfO2 film toward the electrodes. Due to their tetragonal nature, these regions consist of nonswitch-ing (from the ferroelectric point of view) transitional material *(*T-HfO2*)* [55,60,61]. Therefore, the complete device stack consists of TiO*x*Ny*/*TM-HfOx*/*FE−HfO2*/*TM-HfOx*/*TiO*x* sandwiched between two TiN electrodes (see Fig. 8 for details). Further this material intermixing that results in formation of a thin ∼1 nm thick sub-stoichiometric TiO*x* and TiO*x*N*y* regions may affect the effective *k*-value of the stack. The interfacial regions are characterized by a large defect density (oxygen vacancies; *V*o). According to the TEM study, these sub-stoichiometric low-*k* TiO*x* regions are accompanied by a roughly 1*.*5 ± 1 nm thick HfO2 interface characterized with a higher *k*-value due to the presence of

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measured on the bare HfO2 oxide film after top electrode removal showed that a specific device comprises a random distribution of grain sizes. Moreover, the total area of the investigated capacitor (33000µm2*)* consists of hundreds of thousands of grains which all might have a slightly different domain orientation, coercive field or remnant polarization. To address the polycrystalline nature of the film, the modeled hafnia ferroelectric layer is divided into randomly distributed grains and two interface layers next to the electrodes. The granular morphology and grain boundaries play very impor-tant roles since they behave as the preferable locations for the accumulation of the *V*O-s. In order to model the variability in the resulting switching characteristics, each of the modeled grains should represents an averaged ensemble of domains with similar properties. Each ensemble is defined by a dis-tinct coercive field, remanent polarization value and *k*-value. The coercive fields used for the ensembles can be chosen based on the switching densities observed in the First-Order Reversal Curve (FORC) [62] distributions (see Fig. 9b). In Ref. [63,64] it was reported that grains can be correlated with discrete ferroelectric domains and switching events within highly scaled FeFETs (details and modeling approach will

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be discussed in Sect. 4). Further to address the phase transi-tion with field cycling, different phases of the doped hafnia, which can coexist inside the same grain: (a) the orthorhombic phase, responsible for the ferroelectric switching; (b) mon-oclinic, which are not active from the ferroelectric point of view and (c) tetragonal phases, responsible for AFE behavior should be taken into account [41,55].

To account for the behavior of a ferroelectric, the model has to take the history-dependent charge-voltage relation-ship of the ferroelectric [55,65,66] into account. Ferro-electric behavior can typically be simulated using either a Preisach model of hysteresis or the Landau–Khalatnikov the-ory of phase transitions. The differences between those two approaches and the need for the L–K approach in case of ultimately scaled devices will be discussed later in Sects. 4.2 and 4.3. The hysteretic behavior of the ferroelectric is most

where *Q* is the charge of the diffusing space, *λ* is the jump dis-tance, *E*A*,*D istheactivationenergyforion/vacancydiffusion, and *E*eff the effective field along the jump direction, *E*A*,*D the activation energy (energy barrier) given by *E*A*,*D = 0*.*7 eV and *E*A*,*D = 1*.*5 eV for O2-ions and positive *V*O-s [71], respectively. *E*A*,*R represents the activation energy for recombination between complementary species.

In the same study [55,59,72], recombination and diffusion processes were coupled to the charge transport model.

At the end, to emulate the stress-induced defect genera-tion, the thermochemical bond breakage model implemented in the MDLab software [69,72,73] has to be considered. Hafnium oxygen (Hf–O) bond breakage generates an oxygen vacancy, which is an electrically active defect (contributes to the TAT transport), and an oxygen ion. The vacancy genera-tion rate G is described by Eq. (6).

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| commonly represented using a Preisach based, numerically | *G ν* exp | � *E*A*,*G − *bE* | � | *,* | (6) |
| stable model of the hysteresis [67] which is readily avail- |
| able within the commercial TCAD packages used for device | = · | − | � |  |  |
| simulations [66] and is governed by: |

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| --- | --- | --- | --- | --- | --- |
| *P* = *c* · *P*s · tanh | � 1 2*E*c | ln*P*s + *P*r *P*s − *P*r | *(E* ± *E*c*)*� | + *P*off*,* | (4) |

where *c* represents the proportionality constant, *P*s the sat-uration polarization, *P*r the remnant polarization, *P*off the offset polarization, *E* the electric field and *E*c the coercive field.

(details of the implementation and model derivation can be found in Refs. [55,65–67]).

In the next step, to account for a parasitic influence of the dielectric leakage on the ferroelectric switching a Nasyrov-type multiphonon mediated trap-assisted-tunneling (TAT) model was implemented [55,66,68]. Every high-*k* material contains a significant number of defects such as dislocations and impurities which act as electron traps. To take those trap states into account they are modeled as single level acceptor and donor traps within the TCAD software.

These oxygen vacancies actively participate in the TAT current and are responsible for the local field modifications due to charge trapping.

Further, to access the possible material diffusion due to the application of high electric fields, the diffusion and recom-bination of oxygen ions and vacancies have to be calculated through a kinetic Monte Carlo (kMC) model implemented in the MDLab package [69,70]. Recombination rate *R*R and diffusion rate *R*D can be calculated using the following equa-tions:

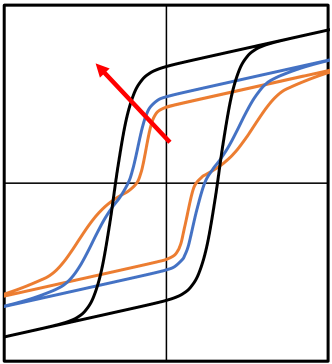
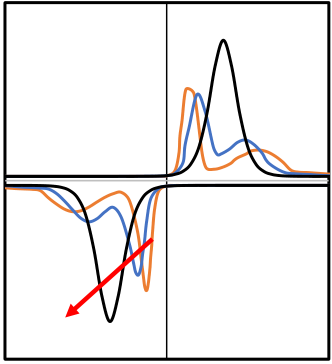
where *ν* is the bond vibration frequency, b the bond polariza-tion factor, *E* the electric field, *E*A*,*G the activation energy for generation of an ion-vacancy pair, kB the Boltzmann’constant and *T* the temperature.

**3.1 Internal bias field and wake-up**

In Ref. [62] it was reported that pristine double current peak behavior could originate from different internal bias fields within the device stack. In the same study it was assumed that different local internal biases or a different effective field manifestations on certain grains and even portions of the device may occur due to a nonuniform defect distribu-tion, polycrystallinity of the annealed film and consequent *k*-value nonuniformity [55,65]. As reported in the previous section, the origin of the observed wake-up in the ferroelec-tric memory is attributed to the progressive decrease in the local internal bias fields. Peši´c et al. reported that the root cause for this decrease is an *V*O defect redistribution that can change their charge state, and/or induce partial phase transi-tions and *k*-value changes within the layer [55]. Within this study defect evolution with cycling was studied by monitor-ing the static leakage current in different lifetime stages of the device. It is very important to note that the density of defects evaluated from the leakage current corresponds mainly to those located at the grain boundaries, where they account for the main contribution to the leakage current [69]. More precisely, the leakage current through the grain boundaries is

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| *R*D = *ν* · exp | �−*E*A*,*D − *Q ~~λ~~* ~~2~~*E*eff | | | � | ; | (5a) | more than tenfold higher than through the grain itself. Thus, |
| the diffusion of the vacancies though the grain would keep |
| the leakage current constant during the wake-up stage. The |
| *R*R = *ν* · exp | �−*E*A*,*R | � | *,* | (5b) | role of O vacancy/ion diffusion is also consistent with the |
| fact that HfO2 based devices are known for their high oxy- |

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| gen mobility [57]. This claim is strengthened by the fact that the electric field applied to HfO2-based ferroelectrics is at least one order of magnitude higher than the fields typically applied to PZT-based films [57]. Furthermore, a recent trans-mission electron microscopy study confirmed the generation and movement of oxygen ions and vacancies within 10 nm thick HfO2 film under similar operating conditions [74]. Starschich et al. proved resistive and ferroelectric switch-ing operation in the same *Y*: HfO2-based device [75]. It is widely accepted that oxygen vacancy motion is required for resistive switching in HfO2 [70].  To address the influence of cycling and dynamics within the stack as well as its influence on the field distribu- | **(a)** | **(b)** | |
| Current [a.u.] | Polarization [a.u.] | **Wake-up** |
| **Wake-up** | | |
| Voltage [a.u.] | Voltage [a.u.] | |
| **Fig. 10** Simulated wake-up of the ferroelectric MÌM capacitor com-prising pristine, intermediate and woken-up **a** current–voltage and **b** polarization–voltage characteristics | | |

tion (predominantly on internal bias fields), the diffusion   
and recombination of oxygen ions and vacancies through ting the grains to a low-*k* (with respect to the orthorhombic a kinetic Monte Carlo model implemented in the MDLab grains), dielectric state.

package [70,76] was simulated. Diffusion rates (Eq. 5a) were calculated by considering activation energies for ion and vacancy diffusion of 0.7 and 1.1 eV, respectively [74,77]. Based on the previous discussion, the constellation of the pristine state within the model was set so that the transitional regions at the electrode interface have much higher defect density. Moreover, based on a TEM study [55] the pristine condition (and internal bias) was generated considering the nonswitching interfaces which possess different *k*-value with respect to the switching fractions of the film which conse-quently affects the distribution of the local fields. In addition to the low-*k* value this additional local internal bias field com-ponent was created by highest density of the vacancies in the interfacial regions [55,65].

This scenario was used to simulate the preferable move-ment of the O vacancies driven by the applied electric field. Simulation of bipolar stress cycling revealed that O vacancies redistribute uniformly within the grain [55]. In addition to this drift/diffusion process, recombination can occur. Due to the recombination of vacancies and interstitial ionstheinternalbiasfielddecreases.Asaresult,uniformfield distribution is created within the stack, resulting in homo-

Using the Preisach model [67], a double peak current–voltage characteristic as well as a pinched hysteresis loop was successfully simulated (see Fig. 10). In addition domain de-pinning was represented by removing the charges previ-ously located at the electrode interfaces, whereas the phase transformation was included by changing of the respective *k*-values and setting the portions of interface as well as the previously nonswitching grains to a ferroelectric active state. These modifications resulted in merging of the current peaks or in other words the opening of the hysteresis curve (Fig. 10), that completely emulates the behavior of the real device.

It should be noted that even the film is woken-up, due to the polycrystalline nature of the film, a slight deviation of experimentally determined *P*r from the theoretical, cal-culated remanent polarization were usually reported. On the other side the remanent polarization calculated for hafnia and zirconia from various authors and from [78] assuming dif-ferent unit cell sizes from doping with values ranging around

Assuming isotropic orientation of grain in a polycrystalline *P*calc = 50 µC*/*cm2and a single polarization direction.

film, the expected average remanent polarization is

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| geneous switching of all domains within the device [55]. | *P*r =*π* 8*P*calc ≈ 21 µC cm2 *,* | (7) |
| Temperature accelerated wake-up experiments reported in |  |
| [55] motivated a simulation [55] which consistently pointed | which matches experimental values very well. |
| out that increase in temperature, results in facilitated vacancy |

diffusion. Hence, yielding more mobile vacancies and faster   
achievement of a uniform electric field within the ferroelec- **3.2 Fatigue** tric [55].

In the same study, it was reported that field driven vacancy distribution during the wake-up stage increase the leak-age current component through the grains, however this is insignificant compared to the total current governed by leak-age thought the grain boundaries.

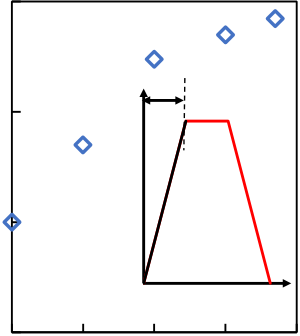
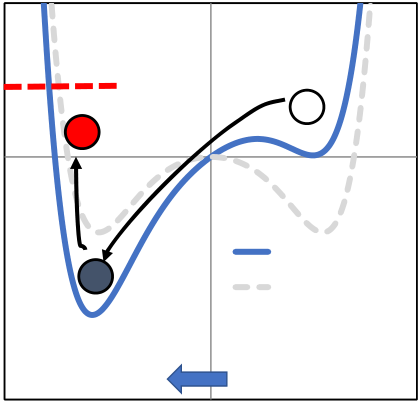
The field cycling endurance of a ferroelectric capacitor may be limited by two aspects: (a) degradation of the dielectric which eventually leads to a hard breakdown and (b) reduc-tion of the switched polarization leading to a closure of the ferroelectric memory window. The later aspect is typically

The interfacial regions are considered nonswitching in the referred to as fatigue.

pristine state. Hence, they act as a passive (dead) layer [55]. The monoclinic (nonswitching) phase was modeled by set-

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First, we want to focus on the time dependent dielec-tric breakdown (TDDB). Detailed comparisons of the DC



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| and AC TDDB lifetime of the ferroelectric capacitors was | **(a)** | | | **(b)**   1E+7 | | | | | |
| reported by Masuduzzaman et al. [79,80]. In these reliability | **EB** | **Bond break**  **energy** | |
| studies, it was shown that the traditional breakdown theories | Endurance | 1E+6 | 1 | Amplitude [a.u.] | **tr**  Time [a.u.] | |
| Free energy [a.u.] | **Overshoot** | **Initial state** |
| for gate oxides are not sufficient to explain the AC TDDB |
| stress dependence of ferroelectric materials. Devices under |
| test comprised a 70-nm thin ferroelectric PZT film. Similar | w/ El.field | 1E+5 |
| to the standard (nonferroelectric) oxides DC TDDB lifetime |
| w/o El.field |
| *(T*BD*)* followed a power law with increasing voltage, char- | **New state** | Electric field | 1E+4   0.1 |
| acterized with a rather small Weibull slope (*β*DC ∼ 1*.*5). Moreover, employing the conductive PFM technique it was | 10 | | 100 1000 |
| Polarization [a.u.] | | | Pulse rise/fall time [us] | | | | | |

shown that the breakdown (BD) spot was always located within the grain boundaries. However, when transferring to AC stress, authors observed drastic decrease in the *T*BD fol-lowed with significant change of the Weibull slopes. The Weibull slope *β*AC for AC stress was significantly increased∼9.1. The reason for this behavior can be explained by mod-eling of hot ion (called hot atom by the authors of [79,90]) degradation which indicated that the increased Weibull slope originates from the breakdown occurring through the grains in contrast to the previously discussed DC induced break-down taking place at the grain boundaries.

The same authors reported a hot ion overshoot to be responsible for the degradation and bond breakage due to alternating voltage stress and subsequent ion crossing of the barrierwithinthedouble-wellpotential(knownfromthermo-dynamic Landau theory) describing ferroelectric materials. Equation (6) describing bond breakage can be modified including the local polarization *p* in such way that the bond breakage energy is represented as

**Fig. 11** Hot ion degradation concept. **a** Double-well energy potential for each switching event, **b** breakdown-endurance dependence on the rise/fall time of the stress pulse. (*inset*) Pulse with variable rising edge

to this overshoot). This bond breakage results in the gen-eration of the defects, i.e., *V*O-s and interstitial O ions in this case, which finally lead to dielectric breakdown. Simu-lations by Masuduzzaman and coworkers [79,80] show that with the increase in the length for the rising/falling edge a strong decrease in the overshoot and corresponding increase in the lifetime of the device was obtained. In order to examine the applicability of the concept, endurance experiments were performed on the HfO2-based MIM capacitors. The plateau width of the pulses was kept constant (1 µs) whereas the ris-ing/falling time of the stress pulses were varied from 100 ns to 1 ms (Fig. 11b, inset). Indeed, the increase in the rise/fall time of the stress pulses resulted in improved endurances. Even though the experiment was successfully repeated, a detailed statistical study is needed in order to draw a final conclusion

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| *E*A*,*G = *E*A*,*G − *pE,* | (8) | about the applicability of the method. |
| Concerning the switching induced degradation, in study |

Since polarization is a dynamic parameter it is described with the following differential equation

by Pesic et al. [59] endurance of HfO2-based ferroelec-tric MIM capacitor devices recorded while stressing with unipolar pulses and compared to endurance monitored while

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| *A*d2 d*t*2 *(p)* + *B* d d*t(p)* − *F (p)* = *E (t)* | (9) | stressing with bipolar pulses. Similar to the results of the |
| breakdown study by Masuduzzaman et al. [79,80] it was |
| reported that only the alternating switching, i.e., a continu- |

where *A* and *B* are inertial and damping coefficients, respec-tively, and *F(p)* denotes internal field originating from double-well ferroelectric potential and corresponding Lan-dau coefficients (*α*, *β*, *γ )* [79,80] and is described with

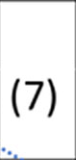
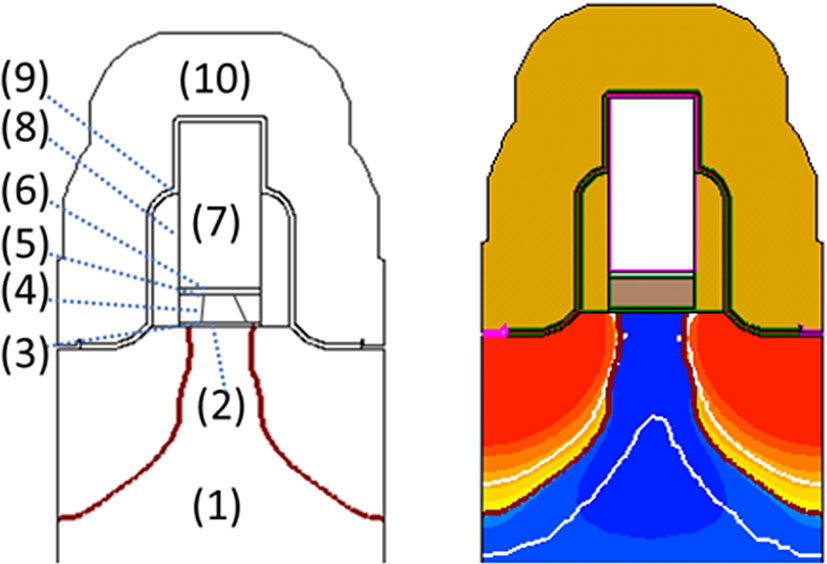
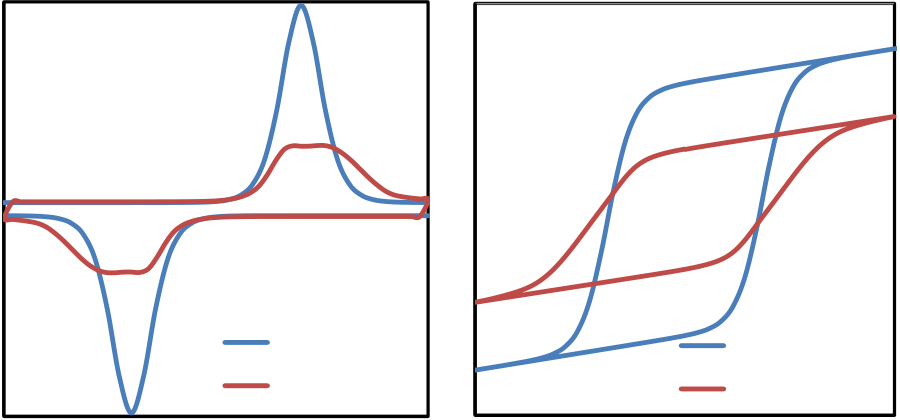
ous change of the polarization state, results in a degradation and consequent reduction of the MW. In contrast the unipo-lar stress does not influence the MW significantly [59,81,82]. Leakagecurrentdefectspectroscopyprovedthatindependent

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| equation | |  |  |  | | --- | --- | --- | | �*α* | *p*2 *β* | *p*4 *γ p* | | � | (10) | of the polarity of the unipolar stress pulses, both leakage cur- |
| *F (p)*d | rent and memory window stayed constant. Analogues to the |
| hot ion degradation model, it was concluded that alternating |
| = −d*t* | 2 + 4 + ∇ |  |  | polarization switching itself (continuous ion displacement |
| within the ferroelectric crystal lattice) induces the endurance |

Combining Eqs. (6)–(10) it was shown that an overshoot of the ion can be caused if pulses with ultra-steep edges are applied. During the change of the polarization state, the ion has a kinetic energy high enough, not just to overcome the barrier but also to climb to the opposite wall of the energy potential eventually reaching the bond breakage energy (due

degradation and fatigue. A direct correlation between the increase in the defect concentration extracted from the charge transport model of the ferroelectric capacitor and the degra-dation of the MW during the fatigue stage was reported [59]. Beside the degradation of the bulk of the device it was reported that faster O vacancy defect generation occurs

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| **(a)**  **(c)**  **Current [a.u.]**  Woken-up  Degraded  **Voltage [a.u.]** | **(b)** |
| |  |  |  |  | | --- | --- | --- | --- | | **Polarization [a.u.]** | **(d)** | Woken-up | **Fig. 13** 2D geometry of the multi grain FeFET device. **a** FeFET geom- | | etry consisting of the Si substrate (*1*), SiO2 interface buffer layer (*2*), | | bottom interface layer (*3*), Si: HfO2 layer (*4*), top interface layer (*5*), | | TiN metal gate (*6*), poly Si (*7*), SiN spacer (*8*), SiO2 liner (*9*) and nitride layer (*10*). **b** Doping profile of the simulated FeFET | | b. Pure electrostatic influence of shielding (charge trapping | | Degraded | increases field over the interface and decreases over the fer- | | roelectric), is accompanied with the domain pining. The | | **Voltage [a.u.]** | |

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| **Fig. 12** Charge trapping influence on the electric field: polarization response and corresponding domain pinning due to the charge trap-ping of **a** woken-up and **b** fatigued stack. For the reasons of clarity single-domain pinning is illustrated. Interface, IF comprises parasiti-cally grown TiO*x* as well as nonswitching portion of HfO2. Comparison of the woken-up and degraded state obtained by simulation: **c** current and voltage and corresponding **d** polarization and voltage response | trapped charge creates dipoles impeding the switching of the domains, which results in a partial or even complete pinning of the domains [59]. For simplicity, the simulated example given with Fig. 12 shows charge trapping within one corner of the top interface region, which alters the local field dis-tribution and pins the domain in the current grain (bounded with the dashed cylinder). As in the measurements [55], also |

the simulated current–voltage traces resulted in broadened

|  |  |
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| within the TiO*x* interface next to the electrodes [59]. Since the bond breakage heavily depends on the applied field and | peaks with decreasing magnitude. As consequence of the altered field, the simulated polarization hysteresis was char- |

the activation energy, usually given by the stoichiometry of the film, these trends can be concluded from Eq. (6) describ-ing the thermochemical bond breakage.

These sub-stoichiometric, nonswitching interface regions characterized by a high number of defects and a lower permit-tivity resulting in higher field drop, yield an increased factor of degradation [65] and result in faster bond breakage. The subsequent charge trapping on this generated defects signifi-cantly influences the local field distribution across the device stack. The field over the interface increases, while it reduces inside the ferroelectric bulk [59,65]. As a consequence, the ferroelectric layer (switching/active part of the device) expe-riences a lower field, which leads to a reduced number of switching domains and decrease in the MW.

In Refs. [55,59] the modeling of the degradation and its influence on the ferroelectric switching was performed. Extracted defect concentration and obtained defect distribu-tion were used as input for the 3D grain boundary model of the MIM capacitor reported in [55] and discussed in Sect. 3.1. The degrading influence on the field distribution across the stack and the resulting change of the current–voltage and

acterized by smoother transitions and lower *P*r (see Fig. 12c, d).

**4 Toward ultimately scaled 1T-memory**  **architectures**

In this section an overview of the current progress in model-ing approaches utilized in investigation of ultimately scaled hafnia-based 1T architectures is presented. Even though var-ious studies reported modeling of the memory window of FeFETs, rather few addressed the limitations caused by switching kinetics and nucleation of the domains in ulti-mately scaled devices. Exactly these properties together with multigrain operation and multilevel-cell (MLC) possibilities will be addressed. Moreover, an overview of the progress in understanding on the neighboring cell disturb through the mix-mode simulations will be reviewed. For this purpose, a 2D multi grain model of the FeFET device was utilized (see Fig. 13). Details about the model development can be found

polarization–voltage characteristic is depicted in Fig. 12a, in [71,83].

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| **4.1 Switching kinetics–nucleation-limited model**  The polarization reversal in the ferroelectric films is influ-enced by various parameters. Among others these are struc-tural and geometric properties of the film as well as electric, thermal and mechanic excitation parameters. For instance, the polycrystalline nature of the film (see Sect. 3 for details), the duration and amplitude of the applied electric field, and the temperature are some of the most important ones. The switching kinetics has both theoretical and practical rele-vance, because it directly impacts the performance of the device containing the ferroelectric.  Many modeling approaches concerning the switching kinetics have been proposed in order to explain the domain nucleation and growth within the material and to relate these processes to the macroscopic electrical manifesta-tions. Two models are generally accepted in this regard. The Kolmogorov–Avrami–Ishibashi (KAI) model is based on the nucleation and propagation of reversed domains [84]. Here, the coercive voltage *V*C, obtained as the intersection between the *P*–*V* hysteretic curve and the voltage axis, follows the simple power law relationship *V*C ∝ *fβ*, where *f* is the exci-tationsignalfrequencyand*β* thefittingparameter.Themodel has been successfully applied for bulk crystals and clean | **Fig. 14** Field and time dependence of ferroelectric switching in ultra-scaled FeFETs. **a** Gate voltage waveform with logarithmically increasing pulse width *t*PW. After each pulse a fast readout of a transfer curve is carried out. **b** *V*T versus *t*PW graph corresponding to 20 repeti-tions of procedure in (**a**), shown for four different VP levels. **c** Standard deviation versus mean value of switching time *t*SW extracted from (**b**).  **d** Fitting of switching probability curves shown in (**b**) |

epitaxial films [85]. On the other hand, polycrystalline disor-  
dered ferroelectrics strongly deviate from this relation [86]

and therefore, several domain nucleation switching limiting (NLS) models have been proposed [86,87]. In these NLS models, the time of domain wall movement is neglected com-pared to the nucleation time. A particularly suitable model for describing field and temperature dependent polarization switching is the one proposed by Du and Chen [87]. Here, the nucleating domains must overcome a critical size defined by an attractive potential. Following a similar approach of Merz [88] for the free energy of domain nucleation within the framework of the classical nucleation theory, a simple expression for the average nucleation waiting time can be

switching kinetics by applying proper programming pulses at the gate electrode, and assessing the resulting polarization state by determining the threshold voltage *V*T of the transis-tor. To this purpose, the gate voltage pulse sequence shown in Fig. 14a is adopted. The *V*N pulse sets the device in the high-*V*T state and with the following *V*P pulse an attempt is done to reverse the polarization and set the low-*V*T state. Due to the fact that the switching of single domains is a stochastic process when they are excited in the proximity of their coercive voltage, the experiment is repeated 20 times in order to collect the statistical set of data. As a result, the

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| found [64]: | � *α* 1 | � | (11) | switching time for a fixed *V*P spreads over approximately |
| *τ τ* exp | one decade, and its mean value decreases with increasing |
| *V*P (Fig. 14b). Reporting the experimental standard devia- |
| =0 · | *k*B*T*·*V*2 |  |  | tion *σt*SW as a function of the average switching time ⟨*t*SW⟩in a log-log graph (Fig. 14c), a slope of unity over several |

where*τ*0 istheshortestnucleationtimepossible,*α* isaparam-eter related to intrinsic material properties including domain wall energy and the portion of a switched polarization by the nucleus, *kB* is the Boltzmann’ constant, *T* is the temperature, and *V* is the voltage necessary to switch the polarization.

In order to experimentally study the switching behav-

decades can be clearly identified. This strongly hints at the stochastic switching governed by a Poisson process.

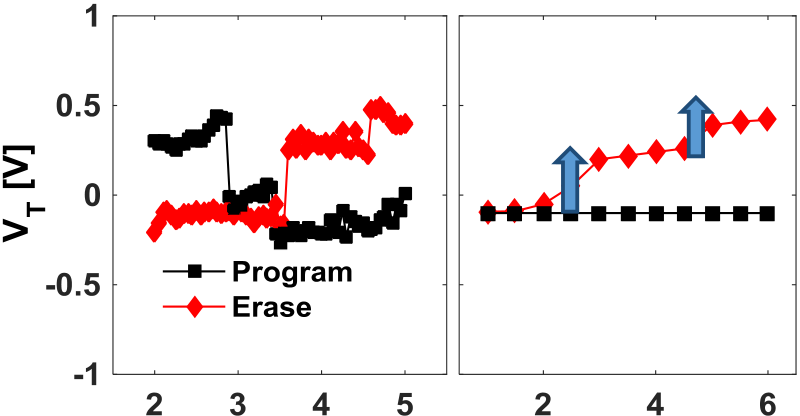
In following the switching in thin films will be modeled and discussed using a purely nucleation-limited approach. Given the small size of grains (the mean grain radius is 10 nm), it is a reasonable assumption that they contain only

ior of the ferroelectric hafnium oxide at the domain level, one domain [89].

ultra-scaled FeFET were characterized. Due to the ultimately reduced device size, the gate stack contains only one or a few ferroelectric domains, which control the channel conductiv-ity [1,19]. In this way, it is relatively easy to monitor their

We start by supposing that upon the application of the electric field the switching is initiated when a relatively small number of critical nuclei are generated within one grain. The generation rate is given by *λ* = 1*/τ* in Eq. (11). Within this

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| picture, the polarization of the grain is considered reversed when a certain critical number *n* of generated nuclei merge | **(a) measured (b) simulated** |

together into a single domain occupying the entire grain.   
Assuming the nucleation to be the Poisson process, the time   
elapsing between each critical nucleus generation *�T*i will   
be exponentially distributed as

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| *p�Ti* = *λ e*−*λ �Ti* | (12) |  |  |
| where *p�Ti* is the probability density function of *�T*i . Thus the overall domain switching time will be given by the sum  of *n* individual *�T*i intervals corresponding to the number of  critical nuclei necessary to form the domain: | |
| **Fig. 15 a** Discrete switching events for both PRG (*black*) and ERS (*red*) of the cell. **b** Simulation of the discrete switching based on the 2D multigrain TCAD model of FeFET (Color figure online) | |

*t*SW =�*n i*=1*�Ti*  (13)

with the mean and variance given by Eqs. (14) and (15),

respectively:

(Fig. 15) was reported. To study the switching to the low *V*T state the measurement was carried out as following: after initial 100 bipolar cycles used for device preconditioning

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| *< t*SW *>*=*n*  *σ*2 tSW= *n λ*2 | (14) | and reach of the fully woken-up state, a reference ERS pulse |
| defining the high reference *V*T was applied. After the read- |
| (15) | out of the reference *V*T, the PRG pulse was incrementally |
| increased in 50 mV steps. It should be noted that every ERS |
| and PRG operation was followed by a readout. Vice versa to |

Now, it is straightforward to determine *n* and *λ* from the experimental data for ⟨*t*SW⟩ and *σt*SWusing Eqs. (14) and (15). Generating then *n* values of exponentially distributed*�T*i as dictated by Eq. (12), summing them up according to Eq. (13) and repeating this procedure 20 times for each pulse width *(t*PW*)* as in the experiment, it is possible to sim-ulate the probabilistic switching. As shown in Fig. 14d, the switching probability curves, extracted from Fig. 14b, are well fitted across the whole time range, with *n* = 5 nuclei for the considered device.

Moreover, the experiment shown in Fig. 14 testifies a clear bias-time trade-off for this nucleation-limited switching [64]. Indeed, referring to Fig. 14b, the device can be programmed at *V*P as low as 2.2 V, but only if *t*PW is longer than 100 µs. However, *t*PW decreases by more than a factor of 100 when *V*P is increased by only 0.6 V. Such exponential dependence, expressed by Eq. (11), represents a source of flexibility and of new opportunities for a future memory design.

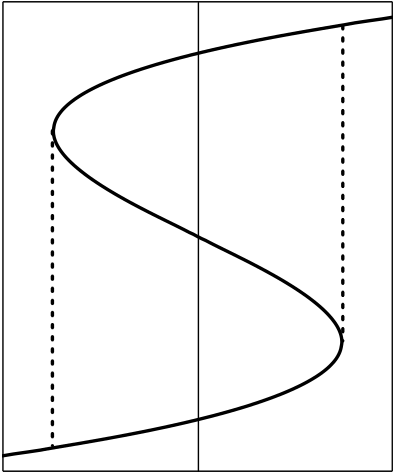
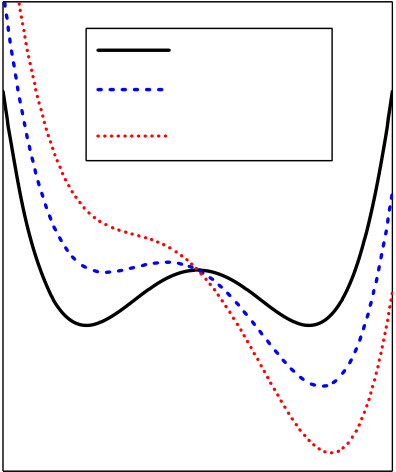
**4.2 Single-domain switching modeling**

As mentioned in the introduction, discovery of the ferro-electric properties in hafnia enabled a tremendous decrease in the memory cell size and high densities of the ferroelec-tric memories. Reasonably, as a next step beyond further geometrical scaling, multilevel-cell operation could result in much increased memory densities per chip area. In 2015 in Ref. [63] a single-domain switching in ultra-scaled Si:HfO2-based FeFET which results in a stepwise change in the *V*T-characteristics for both program and erase sequence

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PRG sweep, pulse sequence was applied for the ERS sweep (red characteristics of the Fig. 15b). Considering an aver-age grain diameter in the order of 10 nm, these ultra-scaled devices (gate length = 28 nm and width = 80 nm) are expected to contain about two to four ferroelectric grains. Further scal-ing of the device while preserving the grain/domain size would result in increased variability of the MW and hence might influence the behavior of the whole array. However, different technological measures can be adopted to coun-teract that issue by decreasing grain size or increasing the uniformity of the ferroelectric film properties.

Here, each discrete switching event which abruptly shifts the *V*T isrepresentedwithadiscretestepin *V*T characteristics. In the same study [63] a simulation was performed to ver-ify the model and physical process behind discrete switching events. Details of the model were reported in [71]. A three-grain 2D TCAD model (see Fig. 13) was utilized for the simulation of the discrete switching. Each grain is charac-terized by a different *E*C value (1, 1.5 and 2 MV/cm). The simulation results for the 2D multigrain TCAD model and its comparison with measurements are shown in Fig. 15. Simi-lar to the measured characteristics, steps in simulated *V*T-*V*G dependence are visible. However, in contrast to the mea-sured abrupt switching steps, the simulated device exhibited smeared out switching steps. Such behavior is a conse-quence of the nature of the Preisach-based model provided in TCAD as discussed in Sect. 3. In contrast to Landau–Khalatnikov model characterized with abrupt switching, the Preisach model which sums up discrete square-shaped hys-terons and averages it over the parameter distribution (e.g.,



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| internal bias fields and coercive fields). It should be noted that | **(a)** | E = 0  0 < E < EC | **(b)** | +Pr |  |
| even within the hysteresis of the three single grains, partial |
| switching and sub-loops are allowed, consequently yielding | Free Energy (a.u.) | Polarization (a.u.) | +EC |
| E > EC |
| a gradual shift of the *V*T. Further, it can be debated if this |
| is physically justified and how small a single hysteron is. It | Polarization (a.u.) | -EC |
| could represent a single unit cell or also a single grain. The |
| results found in this studies, suggest that the latter assumption |
| is more appropriate. Therefore, for ultimately scaled devices | -Pr |
| comprising just a few grains, the usage of a Preisach model |
| that accounts for a large ensemble of domains is no longer |
| Electric Field (a.u.) |
| suitable. Instead, it can be concluded that for the purposes of |

the simulation of discrete switching events and addressing of the single-domain switching kinetics a Landau–Khalatnikov (L–K, see Sect. 4.3 for details) model for emulation of fer-roelectrics is necessary. Detail about the L–K model and its advantages will be addressed in following section.

**Fig. 16 a** Landau free energy potential as a function of the applied field *E*. **b** Polarization–electric field hysteresis calculated from (**a**)

(LK) equation is applied [92]:

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| **4.3 Landau–Khalatnikov model** | *ρ ∂ P∂t*= − *∂u∂ P,* | (17) |
| Starting purely from symmetry considerations, Landau the- |  |

ory is a powerful tool for describing a multitude of phase transition phenomena based on thermodynamic free energy potentials [90]. This phenomenological approach was first applied by Devonshire to model phase transitions in ferro-electricmaterials [91]. Closetothetransitiontemperature *T*C, a high symmetry paraelectric phase transforms into a lower symmetry ferroelectric phase. The simplest way to describe this phase transition is by doing a series expansion of the free energy *u* using the order parameter *P*, which is the electrical polarization:

*u* = *αP*2+ *β P*4+ *γ P*6− *E P.*  (16)

Here, *α*, *β*, *γ* are the ferroelectric anisotropy constants and *E* is the electric field in the ferroelectric. Below *T*C, for second-order phase transitions, *α <* 0, *β >* 0 and *γ* = 0, whereas for first-order phase transitions *α <* 0, *β <* 0 and *γ >* 0. Figure 16a shows the free energy as a function of *P* for differ-ent applied fields *E*for a second-order transition. Without an applied field there are two degenerate energy minima corre-sponding to the remanent polarization states (+*P*r and −*P*r*)*. Whenafield *E* isapplied,oneminimumisbecomingenerget-ically more favorable while the other one increases in energy and ultimately vanishes at the coercive field *E*C. By differ-entiating Eq. (16) with respect to *P* and setting d*u/*d*P* = 0, we obtain the *S*-shaped *P*–*E* relationship corresponding to the polarization hysteresis as shown in Fig. 16b. It should be noted here, that in the case shown in Fig. 16b the switching at *E* = *E*C would be instantaneous, which is obviously not physical. Therefore, to calculate more realistic switching dynamics based on the free energy in Eq. (16), the Landau–Khalatnikov

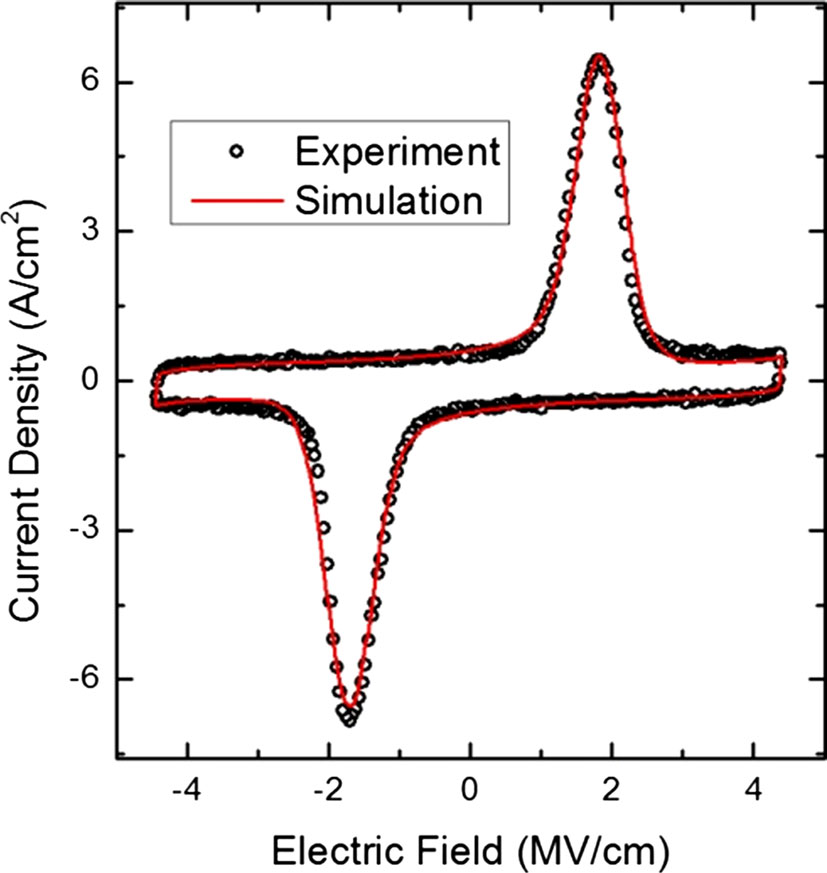
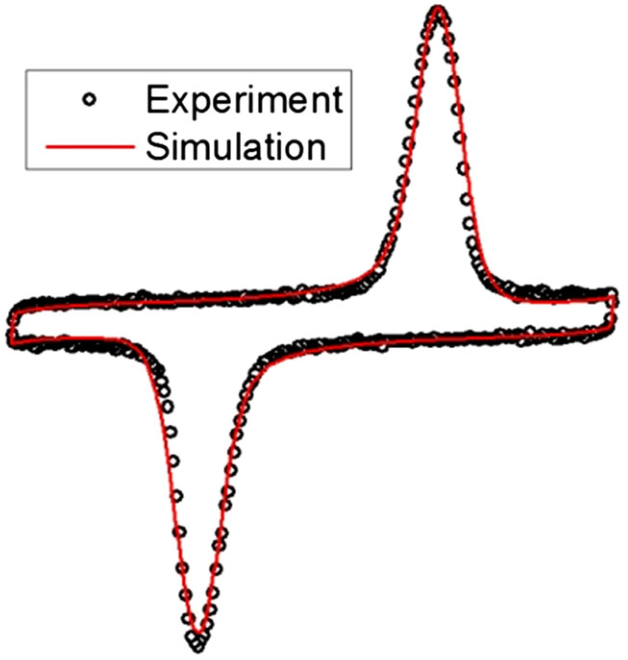
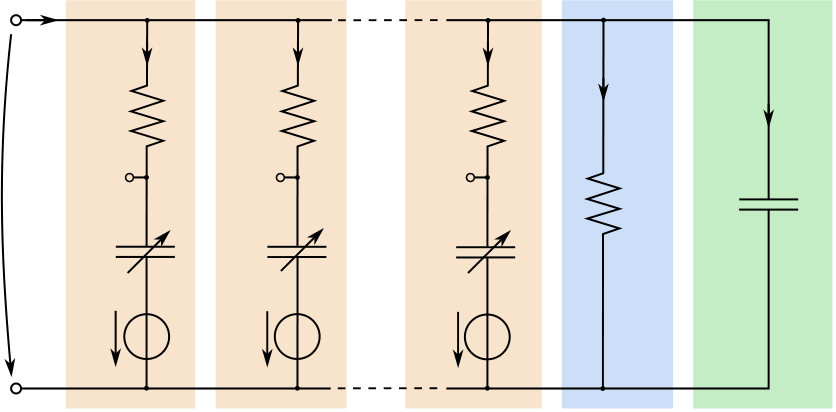
where *ρ* is a damping constant of dimension [*�*m], which corresponds to the internal resistivity of the ferroelectric which limits the change of *P* with time. In case of Fig. 16b, *ρ* would be zero.

It should be noted, that this is a mean-field theory which assumes homogeneous properties throughout the whole fer-roelectric. While this is obviously not true for polycrystalline thin films like HfO2, we will show how inhomogeneities can be incorporated into the Landau–Devonshire model. A gen-eral extension of the model to include a gradient term *(*∇ *P)*2 was proposed by Ginzburg and Landau [93], which is also used in more complex ferroelectric phase field models [94]. However, we will propose a simpler approach to describe HfO2-based ferroelectrics, which is much less computation intensive than phase field models and much more accurate than homogeneous Landau theory.

Generally, LK models based on homogeneous Landau–Devonshire theory are not suited to describe polarization switching in ferroelectric thin films, since local effects like domain nucleation and growth are known to govern polariza-tion reversal [95]. Therefore, a modified LK approach was proposed [96], where the ferroelectric film is partitioned into smaller regions, which are electrically in parallel and are each described by Eqs. (16) and (17). By introducing spatial dis-tributions of the parameters *α*, *β*, *γ* and *ρ*, nonuniformity of the film can be modeled [96].

In the case of HfO2-based ferroelectrics, the polycrys-tallinity and grain structure is especially of interest. Some grains might be paraelectric and therefore can be modeled as a regular linear capacitance in parallel. Each ferroelectric grain, on the other hand, is described by the LK-equation. Then, in case of a ferroelectric capacitor with two metal elec-

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| |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  |  | | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | | |  |  |  |  |  |  | | --- | --- | --- | --- | --- | --- | | iF | Grain 1 | Grain 2 | Grain N | Leakage | Dielectric Grains | | iF1 | iF2 | iFN | iL | iCD | | R1 | R2 | RN | | V | V | V | |  | | |  |  |  |  |  |  |  |  |  | | --- | --- | --- | --- | --- | --- | --- | --- | --- | | VF | i1  (P1) | CF1 | i2  (P2) | CF2 | iN  (PN) | CFN | RL  (VF) | CD | | Vb1 | Vb2 | VbN |   **Fig. 17** Equivalent circuit representation of the multigrain Landau–  Khalatnikov model for a ferroelectric HfO2 capacitor  trodes, the total charge on the capacitor is calculated as  *N*  *N*  (18) |   *Q*FE = *A*FE⎛⎝*ε*0*E* +1�*Pj*⎞⎠ ≈*A*FE�*Pj,*  **Fig. 18** Comparison of experimental data and multigrain Landau–  Khalatnikov simulation of an 18 nm thin Gd: HfO2 capacitor. Current  where *A*FE is the area of the capacitor plates, *ε*0 is the vac- response to a triangular voltage signal with a frequency of 10 kHz  uum permittivity, *N* is the total number of grains and *Pj* is  the polarization of the grain *j*. It is assumed that all grains | |

have the same area *A*FE*/N* and that each grain has a homoge-  
neous polarization. While this is an approximation, it seems   
reasonable since grain sizes in HfO2-based ferroelectrics are (see Sect. 4.2 for details). However, new complications arise

typically in the same order as the film thickness *T*FE (5–30 nm) [97]. Additionally, grain boundaries generally run from one electrode to the other, perpendicular to the film and nucleation-limited switching kinetics show (see Sect. 4.1) that individual parts of the film switch independent of each other [64]. Therefore, the assumption of modeling nonin-teracting grains electrically in parallel seems justified. The

when simulating a FeFET structure due to different boundary conditions compared to a capacitor: When the ferroelectric HfO2 is deposited directly onto the Si channel a SiO2 inter-face forms [64]. Therefore, the potential across each grain will be locally different and a particular spatial distribution of grains will have to be assumed. In case of a regular capac-itor with metal electrodes, the actual spatial position of each

equivalent circuit representation of this model is shown in grain is irrelevant.

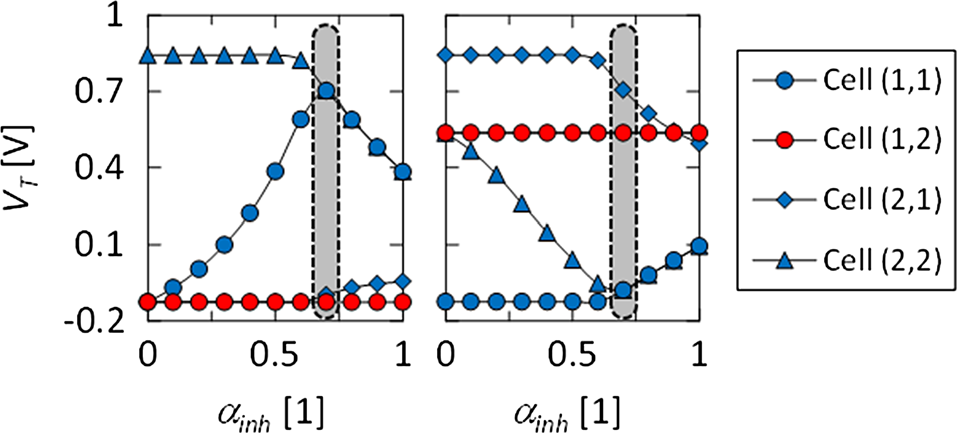
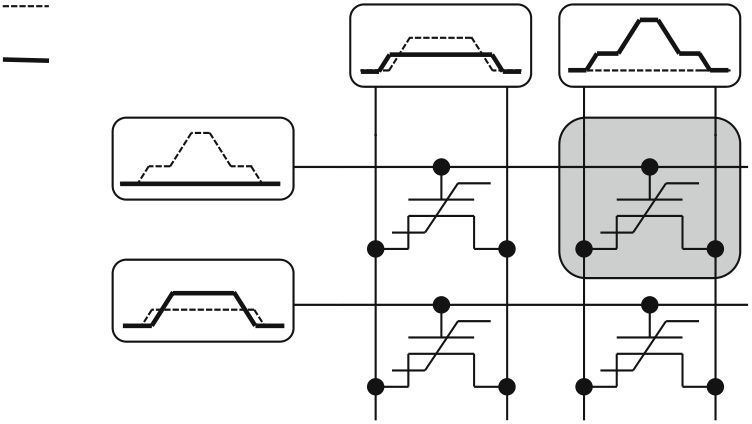
Fig. 17. Here, *R j* = *ρ jT*FE*/A*FE is the internal resistance,

is the internal bias voltage [98] and *i*Fj is the current of grain *Vij* = 2*T*FE*(α j Pj* +2*β j P*3 *j)*+*V*b *j* is the internal voltage, *V*b *j*

*j*. *C*Fj corresponds to the nonlinear capacitance of each grain ferroelectric grain which is given by *(*d2*u/*d*P*2*)*−1*A*FE*/T*FE. The leakage current *i*L is modeled by a parallel resistance *R*L which is a function of *V*F (typically an exponential depen-dence). All nonferroelectric grains can be combined into a parallel capacitance *C*D. Using this multigrain LK approach, experimental data from standard *P*–*V* hysteresis measure-ments can be fitted by adjusting the *α*, *β*, *γ* and *ρ* parameter distributions [96], as shown in Fig. 18 for a 18 nm thin fer-roelectric Gd: HfO2 film, with *N* = 10*,* 000. While this example shows the very good agreement of the multigrain LK model with data for rather large capacitors *(A*FE = 33*,*000 µm2*)*, it should still result in a more accu-rate representation of HfO2-based ferroelectrics in nanoscale devices. For instance, it should be possible to describe the single-domain switching seen in FeFETs scaled to the 28 nm node [64] more accurately compared to the Preisach model

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Additionally, completely new physical phenomena arise when using the LK model compared to other ferroelec-tric models. Based on Landau theory, it was proposed that the region of negative slope of the *S*-shaped *P*–*E* curve (corresponding to a negative capacitance (NC)) could be used to build a FET with subthreshold swing lower than 60 mV/decade at room temperature [99]. Such NCFETs based on Hf1−*x*Zr*x*O2 have already been demonstrated experimentally [100] and are a promising way to further reduce the supply voltage and thus the power dissipation in future logic circuits. However, all device simulations reported so far applied a homogeneous single-domain Lan-dau model [101], which cannot accurately predict effects of nonuniformity in HfO2-based ferroelectrics. The device structure of an NCFET is very similar to a regular FeFET, since also a ferroelectric is used in the gate stack, just with different desired parameters (*T*FE, *P*r and *E*C*)*. Therefore, using an LK-based approach would be beneficial to combine existing models for both memory FeFETs and logic NCFETs into a single comprehensive modeling framework.



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| Program opera�on | SL / BL (n) | SL / BL (n+1) |  |

Erase opera�on

WL (m)

WL (m+1)

|  |  |
| --- | --- |
| **Fig. 19** Program and erase operation of FeFET cells in AND architec-ture unit cell representation. The transient voltage trains on the terminals are illustrated for the program (*dashed lines*) and the erase (*solid lines*) operation. The cell to be written is highlighted in *gray* | **Fig. 20** Verification of *V*DD*/*3 operational scheme by TCAD simu-lation. The evolution of threshold voltages *V*T over the inhibit factor *α*inh is shown for all cells of the AND architecture unit cell representa-tion. The indices of the cells in the legend encode rows/wordlines and columns/bitlines and sourcelines. Cell (1, 2) is the selected cell. *Left* rep-resents the program operation and *right* the erase operation performed |

on cell (1, 2)

**5 Mixed-mode modeling and circuit model**

After the single device simulations discussed up to now, next level in simulation hierarchy is achieved through the grouping of these devices into circuits and performing the mixed-mode simulations. These mixed-mode TCAD simu-lations extend the single device model not only by adding different device models but also by combining it with the HSPICE compact models. Based on that it is not hard to con-clude that these simulations are positioned hierarchy-wise between the device simulations and compact models. In the first part of this chapter a mixed-mode simulation of the AND array architecture of FeFET will be reviewed, whereas the second part is devoted to a circuit model of the FeFET.

**5.1 TCAD simulation of elementary array architectures**

When looking at semiconductor memories the first step to the circuit level requires array-level simulations. This type of simulation is required in order to verify the functionality

cells must not change their binary state. The *V*DD/3 scheme in principal allows for a complete voltage drop of *V*DD across the selected cell whereas the unselected cells only experience one third of this write voltage as parasitic disturb.

In order to verify the functionality of such an operational scheme for a specific FeFET device geometry (see Fig. 13 in Sect. 4), TCAD simulation can be utilized. For the given cell array unit cell of Fig. 19, various worst case scenarios have to be considered (see also [103] for a detailed description). The results of the respective TCAD simulation are shown in Fig. 20.

The TCAD simulations where performed such that all worst case disturb scenarios on the respective cells in the AND architecture unit cell representation were analyzed as a function of the so-called inhibit factor *α*inh. With respect to Fig. 19, the inhibit factor determined the voltages applied to the respective signal lines:   
Program operation:

|  |  |  |
| --- | --- | --- |
| of each individual memory cell when embedded into a larger | *V*WL*(m)* = *V*P  *V*WL*(m*+1*)* =*α*inh· *V*P  *V*S*/*BL*(n)* = *α*inh · *V*P  *V*S*/*BL*(n*+1*)* = 0 | (19) |
| array of multiple cells. For example write disturb is one of |
| the parasitic effects occurring solely when a single cell is | (20) |
| incorporated in a certain memory architecture. In order to | (21) |
| illustrate this, the so-called AND architecture is chosen as an |
| (22) |
| example (see Fig. 19). |
| For FeFET cells arranged in an AND configuration, a gen- |
| Erase operation: |
| eral way to write a selected cell is given by the so-called |
| *V*DD/3 scheme described elsewhere [102,103]. This opera- | *V*WL*(m)* = 0  *V*WL*(m*+1*)* = *α*inh · *V*P *V*S*/*BL*(n)* =*α*inh· *V*P  *V*S*/*BL*(n*+1*)* = *V*P | (23) |
| tional scheme allows for reducing the write disturb occurring |
| in one-transistor (1T) FeFET array architectures. That is, | (24) |
| when a cell is selected to be written (i.e., either for program or | (25) |
| erase operation), the shared terminals called wordline (WL), |
| bitline (BL) and sourceline (SL) have to be raised or lowered | (26) |
| to certain write and inhibit voltages. At the end, the voltages |

have to be chosen such that the voltage drop across the gate stack of the selected cell induces a polarization reversal in the ferroelectric film while at the same time all neighboring

Accordingly, for the program operation in Fig. 20 on the left-hand side it can be observed that only for an inhibit factor of around 0.7, it is possible to reduce the worst case disturb for

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| cell (1, 1) appropriately. To be more specific, the worst case for cell (1, 1), when cell (1, 2) shall be programmed into the low-*V*T state, is present when cell (1, 1) originally resides in a high-*V*T state. This state can only be maintained when the unselected column n is raised to around 0.7 times the write voltage *V*P. This verifies that the *V*DD*/*3 scheme is indeed best suited for minimizing disturb.  The same analysis can be performed for an erase operation oncell(1,2)viatheso-calledPositive–Source–Drain–Erase– |  |

Scheme (PSDES) [103]. According to Fig. 20, also there the *V*DD*/*3 scheme, i.e., at an inhibit factor of around 0.7, works best for reducing write disturb, in this case erase disturb in particular. Especially with respect to the PSDES scheme it is interesting to note that the high-*V*T state after the erase operation on cell (1, 2) is not as high as it is for cell (2, 1) which had previously been written into the reference state by a so-called negative gate erase (NGE) operation. This is rooted in the effectiveness of the PSDES and a detailed computational study can be found in [83]. After a in-depth discussion of the single device behavior and influence of the different cells on the neighboring cells in an array, a circuit

**Fig. 21** CircuitmodelofaferroelectriccapacitorbasedontheLandau–Khalatnikov equation. Ferroelectric capacitor and integrating circuit are marked with *orange* and *blue rectangle*, respectively (Color figure online)

Finally physics-based models are based on the LK-equation [117–119,121] or the KAI-model [120]. The nonlinearity of the ferroelectric is either reproduced abstractly by controlled current and voltage sources [117] or, more circuitry-wise, by a series connection of a nonlinear capacitor *C*N and a resis-tor *R*N with a linear capacitor *C*0 in parallel [118–120]. The linear capacitor originates from the consideration of the total

model will be discussed next. charge in the ferroelectric.

All previously-mentioned models are suitable to repro-

**5.2 Circuit model of ferroelectric devices**

Different circuit models were proposed in the literature to model ferroelectric materials in electron devices. Prior to the actual fabrication step of devices they allow the evaluation of designed circuits. Simulation models comprise mathematics-based approaches where the hysteresis loop is reproduced by mathematical functions, switching current models, the parallel elements method as well as physics-based models (see Sect. 3) based on nucleation mechanisms (KAI and NLS model; see Sect. 4.1) or the double-well potential of the Gibbs free energy (Landau–Khalatnikov model; see Sect. 4.3).

The mathematical approach includes the Preisach model

duce the hysteretic behavior of the ferroelectric. Switching current models can reproduce transient switching and DC behavior, but the switching current has to be known in the first place or assumptions about its behavior have to be made. The parallel elements method, though the single dipoles can be modeled in a very simple way (e.g., Schmitt trigger), can lead to increasing simulation times because of the numer-ous parallel branches. The same approach accounts for the physics-based models when several domains are modeled. The latter ones are favorable when the priority is given to a behavioral model which emulates the physics of the switch-ing kinetics. Static and dynamic behavior can be described and only a low number of parameters is needed to describe

[104–110] already briefly introduced in Sect. 3. Statistically a single dipole.

distributed dipoles with individual coercive fields and a spe-cific density function with respect to the coercive fields are superimposed to obtain the overall polarization. If the density function is Gaussian type, the resulting polarization becomes a tanh-function. In the given example, the full hysteresis loop

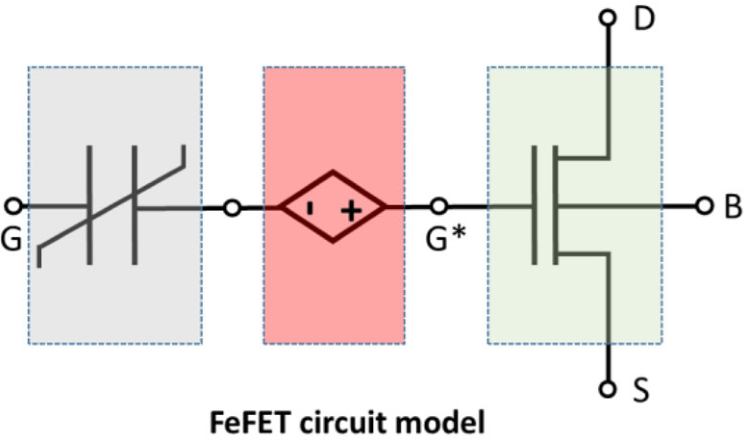
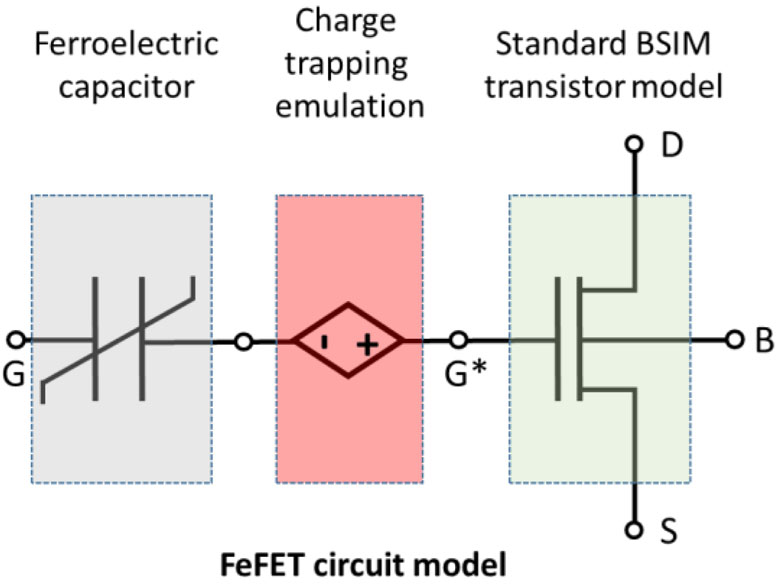
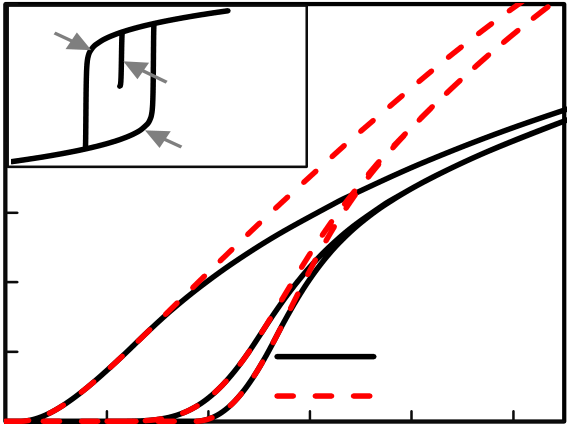
The Model of the ferroelectric material described in the following (Fig. 21) is based on the approach of Aziz et al. [118] who made use of the LK-equation (Eqs. (16)–(17); Sect. 4.3) to express the voltage drop *V*FE across the ferro-electric as:

|  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| is reproduced by two tanh-functions which are shifted along | *V T* | � | *ρ* | d*Q*P | *α Q*P | *β Q*3 | *γ Q*5 | � | (27) |
| the *E*-axis by ±*E*C (see Eq. (4) for details). Another mod- |
| eling approach relates to the polarization switching current | FE =FE |  | *A*FE | d*t* | + | + FE | + FE |  |  |
| of the ferroelectric [111–114]. Here the current is modeled |

based on the rate equations of the switching dipoles [114], precalculated polarization current [113] and first-order relax-ation processes [111,112]. On the other side the parallel elements method [115,116] emulates the structure of a ferro-electric material as a parallel circuit of several ferroelectric domains. Single domains are simulated by a Schmitt trig-ger and voltage controlled capacitor and resistor [115,116].

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where *α*, *β*, *γ* and *ρ* are the parameters of the LK-equation introduced in Sect. 4.3. *T*FE and *A*FE are the thickness and the area of the ferroelectric, respectively. The first term corresponds to a series resistor with the resistance *R*N = *T*FE·*ρ/A*FE and the following terms reflect the nonlinearity of the ferroelectric capacitor. This nonlinearity (of the ferro-electric) is emulated by a voltage controlled voltage source



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**(a)**  ward solution to address this parasitic behavior would be to account this shift by an auxiliary sub-circuit (see Fig. 22a; charge trapping emulation), similar to one reported in [122]. Other implementations like adding a second controlled volt- age source in series are possible. However, since the charge trapping and de-trapping effects occur mainly in the inter- face oxide region and thus depend on the respective electrical fields, the additional circuit should be placed in-between the transistor model and the voltage source modeling the ferro- electric polarization effect.

|  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **(b)** | 1.2E-5 | "1" | 0.0 | unaligned | | Vds = 0.5V | | **6 Summary and outlook** |
| 1.0E-5 | |
| *I*ds [A] | 8.0E-6 | | The discovery of ferroelectric properties within HfO2 revived |
| "0" | |
| the interest in ferroelectric memories leading to a rapid devel- |
| 6.0E-6 | | 0.1 | 0.2 | opment and impressive device scaling in the last 10 years. In |
| 4.0E-6 | | this article, we provided an overview of modeling hierarchies |
| and approaches which were developed in this timeframe not |
| 2.0E-6 | |
| only to unveil the origin of the ferroelectricity in binary |
| Vds = 1V | |
| 0.0E+0  -0.1 | |
| oxides and physical mechanism taking place with field stress- |
| 0.3 | 0.4 |
| ing of the device, but also as strategies that have to be |
| *V*gs [V] | |
| considered when modeling and designing the future ulti- |

**Fig. 22 a** Circuit model of FeFET comprising a FE capacitor, con-trolled voltage source (charge trapping emulation) and standard BSIM model of FET transistor. **b** Results for the on- and off-state of the FeFET from the circuit model shown in Fig. 21. For *V*ds = 0*.*5 V (*solid line*) and *Vds* = 1 V (*dashed line*). The *middle curve* shows the initial curve (dipoles unaligned). The high *V*T-state corresponds to polarization state“0”, the low *V*T−state corresponds to polarization state “1” (see *inset*)

*V*N which is dependent on the charge *Q*P. To get *Q*P, the cur-rent *I*VN through the voltage source is integrated over time by an integrator circuit. Based on this simulation model (Fig. 21) clamped on standard BSIM transistor model (Fig. 22a) typi-cal on-/ off-states of a FeFET are observed (Fig. 22b).

In contrast to the model described here, previously dis-cussedmodel(seeSect.4.3)apartfrommultigrainimplemen-tation, includes internal bias fields and nonlinear capacitance, whereas the model reported by Aziz et al. [118] emulates this nonlinearity with voltage source. Including parallelization (to emulate multidomain nature of the devices) and addi-tional constant voltage source (to emulate internal bias field present) a direct conversion from model described here to model discussed in Sect. 4.3 is possible.

To assess the realistic device performance trapping effects havetobeconsideredandincorporated.Trappingofelectrons and holes as a parasitic effect compensates the MW of the device by shifting the threshold voltage toward more posi-tiveandnegativevalues,respectively.Thisparasiticthreshold voltage shift is proportional to number of defects filled for certain applied pulse amplitude and length. A straight for-

mately scaled devices for practical applications. Even though there is a number of challenges to be overcome, both in the experimental realization and the modeling framework, fer-roelectric memories based on hafnium oxide show a huge potential for becoming a wide-spread, high-speed and low-power nonvolatile memory solution of the future.

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