

A phenomenological cohesive model of ferroelectric fatigue

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Abstract

We develop a phenomenological model of electro-mechanical ferroelectric fatigue based on a ferroelectric cohesive law that couples mechanical displacement and electric-potential discontinuity to mechanical tractions and surface-charge density. The ferroelectric cohesive law exhibits a monotonic envelope and loading–unloading hysteresis. The model is applicable whenever the changes in properties leading to fatigue are localized in one or more planar-like regions, modeled by the cohesive surfaces. We validate the model against experimental data for a simple test configuration consisting of an infinite slab acted upon by an oscillatory voltage differential across the slab and otherwise stress free. The model captures salient features of the experimental record including: the existence of a threshold nominal field for the onset of fatigue; the dependence of the threshold on the applied-field frequency; the dependence of fatigue life on the amplitude of the nominal field; and the dependence of the coercive field on the size of the component, or *size effect*. Our results, although not conclusive, indicate that planar-like regions affected by cycling may lead to the observed fatigue in tetragonal PZT.

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1. Introduction

Ferroelectric materials are extensively used in a variety of sensor and actuator applications, where the coupling between mechanical and electrical fields is of primary interest. They are also a promising set of materials for improved dynamic as well as non-volatile memory devices, where only the electrical properties are directly exploited. However, ferroelectrics are brittle, and their low fracture toughness (in the order of $1 \text{ MPa m}^{1/2}$) makes them susceptible to cracking. In addition, ferroelectric materials exhibit electrical fatigue (loss of switchable polarization) under cyclic electrical loading and, due to the strong electro-mechanical coupling, sometimes mechanical fatigue as

well. Conversely, the propagation of fatigue cracks hinders the performance of the devices and raises serious reliability concerns.

Ferroelectric fatigue is caused by a combination of electrical, mechanical and electro-chemical processes, each of which has been claimed to be responsible for fatigue [1]. Several electro-chemical mechanisms have been posited as the likely cause of polarization fatigue [2], but no general consensus appears to have emerged as yet. Fatigue mechanisms variously include processes of distributed damage over the bulk (see, e.g., Refs. [3,4]) and processes of localized damage, including micro-cracks and ferroelectric–electrode interfaces (see, e.g., Refs. [5–7]). The proposed mechanisms include domain-wall pinning and inhibition of reversed domain nucleation [8,9,2]. The former mechanism is thought to play a dominant role in the bulk, whereas the latter mechanism is thought to operate primarily at or near electrode

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interfaces [10]. The relative roles of these and other mechanisms may depend on the frequency of the applied field [10]. At the atomic level, oxygen vacancies are likely to promote fatigue, e.g., by migrating under the action of the electric field to form extended defects that pin domain walls [11–13]. Purely mechanical mechanisms, such as microcracking, are also likely to play some role. Thus, fatigued ceramic specimens often contain scattered microcracks of size comparable to that of the grains [14,15]. Profuse microcrack clouds have been observed at the specimen edges and surrounding macroscopic cracks [16]. Macroscopic crack patterns are present in some fatigued ceramic specimens [17]. Severe cracking was also observed in barium titanate single crystals subjected to cyclic bipolar electric load [18]. Modeling work suggests that microcracking is indeed a cause of loss of polarization [6].

Experiments on crack propagation have been reported for samples loaded electrically, mechanically, or under combined loading, cyclic or static [19–22]. It is not uncommon for different experiments to lead to apparently contradictory conclusions [23], a testament to the complexity of the phenomenon of ferroelectric fatigue. On the other hand, a large body of experimental data concerns smooth samples, or samples without an initial pre-crack. In these experiments, the main property of interest is the evolution of polarization, and often no special attention is given to the mechanical integrity of the sample. Epitaxially grown thin films constitute a common configuration for tests involving smooth samples [24].

Despite these recent experimental and modeling advances, the precise nature of the interactions between fracture, deformation and defect structures underlying ferroelectric fatigue is in need of further elucidation, and a physics-based multiscale model enabling the prediction of the fatigue life of ferroelectric devices is yet to emerge. Therefore, there remains a need for phenomenological and empirical models that can be experimentally validated and used in engineering design. In this vein, cohesive theories provide an effective means of modeling fatigue-crack nucleation and growth for arbitrary crack and specimen geometries and loading histories [25,26]. As noted by Nguyen et al. [25], the essential feature that a cohesive law must possess in order to model fatigue crack growth is loading–unloading hysteresis. By this simple but essential device, cohesive models of mechanical fatigue have been shown to account for deviations from Paris’s law in metals such as are observed for short cracks and overloads [25]; and to predict fatigue-crack nucleation in smooth-surface metallic specimens [26]. The appealing feature of cohesive models of fatigue is that a single mechanistic, albeit empirical, model applies to nucleation and propagation, short and long cracks, and arbitrary loading histories.

Conventional cohesive models of fracture seek to describe the relation between cohesive tractions and open-

ing displacements at the tip of a crack. Working by analogy, Gao et al. [27–29] have proposed the use of Dugdale-like models for the electric displacement in ferroelectric materials. In this paper, we further extend the concept of cohesive surface to ferroelectric materials by understanding it to be a surface of both *mechanical opening displacement and electric-potential discontinuity*. Correspondingly, we extend the concept of cohesive law to mean a general relation between the work-conjugate pairs: mechanical displacement and electric-potential discontinuity; and mechanical tractions and surface-charge density. Following Nguyen et al. [25], in order to model ferroelectric fatigue we endow the ferroelectric cohesive law with loading–unloading hysteresis. In order to simplify modeling, following [30] we make the additional assumption that the cohesive law derives from a potential that depends on a scalar effective electro-mechanical jump, which is a weighted norm of mechanical opening displacements and electric-potential jump.

Using a simple Smith–Ferrante monotonic envelope and an exponential-decay law of loading–unloading hysteresis, we show that the model is able to qualitatively capture salient features of the experimental record including: the existence of a threshold nominal field for the onset of fatigue; the dependence of the threshold on the applied-field frequency; the dependence of fatigue life on the amplitude of the nominal field; and the dependence of the coercive field on the size of the component, or *size effect*.

2. Electro-mechanical cohesive laws

The essential structure of cohesive laws in ferroelectrics may be elucidated by recourse to a conventional Coleman–Noll argument (cf., [30] for an application to mechanical decohesion). In particular, the pairing between stress-like and deformation-like variables is determined by the internal-power identity or, equivalently, by the virtual-work identity. For a dielectric solid, this identity is [31–33]

$$\int_{\Omega} (\boldsymbol{\sigma} : \delta \boldsymbol{\epsilon} - \mathbf{D} \cdot \delta \mathbf{E}) dV = \int_{\Omega} (\mathbf{b} \cdot \delta \mathbf{u} - \rho \delta \phi) dV + \int_{\partial \Omega} (\mathbf{t} \cdot \delta \mathbf{u} - \omega_f \delta \phi) dS, \quad (1)$$

where Ω is the spatial domain occupied by the solid and $\partial \Omega$ is its boundary; $\boldsymbol{\sigma}$ is the stress;

$$\boldsymbol{\epsilon} = \frac{1}{2}(\nabla \mathbf{u} + \nabla \mathbf{u}^T) \quad (2)$$

is the strain; \mathbf{D} is the electric displacement;

$$\mathbf{E} = -\nabla \phi \quad (3)$$

is the electric field; \mathbf{b} is the body force; \mathbf{u} is the displacement; ρ is the *free* charge density; ϕ is the electric potential; $\mathbf{t} = \boldsymbol{\sigma} \cdot \mathbf{n}$ is the surface traction; and $\omega_f = -\mathbf{D} \cdot \mathbf{n}$ is the *free* surface charge per unit area. For a solid with a surface of discontinuity T , or cohesive surface, we have

$$\begin{aligned}
\delta W_{\text{ext}} &= \int_{\Omega} (\mathbf{b} \cdot \delta \mathbf{u} - \rho \delta \phi) dV + \int_{\partial \Omega} (\mathbf{t} \cdot \delta \mathbf{u} - \omega_f \delta \phi) dS \\
&= \int_{\Omega} (\mathbf{b} \cdot \delta \mathbf{u} - \rho \delta \phi) dV + \int_{\partial \Omega} [(\mathbf{n} \cdot \boldsymbol{\sigma}) \cdot \delta \mathbf{u} + (\mathbf{n} \cdot \mathbf{D}) \delta \phi] dS \\
&= \int_{\Omega} (\mathbf{b} \cdot \delta \mathbf{u} - \rho \delta \phi) dV + \int_{\Omega^{\pm}} [\nabla \cdot (\boldsymbol{\sigma} \cdot \delta \mathbf{u}) \\
&\quad + \nabla \cdot (\mathbf{D} \delta \phi)] dV + \int_{\Gamma} [(\mathbf{n} \cdot \boldsymbol{\sigma}) \cdot \llbracket \delta \mathbf{u} \rrbracket + (\mathbf{n} \cdot \mathbf{D}) \llbracket \delta \phi \rrbracket] dS
\end{aligned} \tag{4}$$

and consequently

$$\begin{aligned}
\delta W_{\text{int}} &= \int_{\Omega^{\pm}} (\boldsymbol{\sigma} : \delta \boldsymbol{\epsilon} - \mathbf{D} \cdot \delta \mathbf{E}) dV + \int_{\Gamma} [(\mathbf{n} \cdot \boldsymbol{\sigma}) \cdot \delta \llbracket \mathbf{u} \rrbracket \\
&\quad + (\mathbf{n} \cdot \mathbf{D}) \delta \llbracket \phi \rrbracket] dS.
\end{aligned} \tag{5}$$

For monotonic loading of the cohesive surface, these work-conjugacy relations naturally suggest a free energy per unit surface, or cohesive potential, of the form $\Phi(\llbracket \mathbf{u} \rrbracket, \llbracket \phi \rrbracket, \mathbf{q})$ such that

$$\mathbf{t} = \frac{\partial \Phi}{\partial \llbracket \mathbf{u} \rrbracket}, \tag{6a}$$

$$-\omega_f = \frac{\partial \Phi}{\partial \llbracket \phi \rrbracket}, \tag{6b}$$

where \mathbf{q} is some suitable set of internal variables. A possible additional dependence of Φ on temperature is omitted for notational convenience. Thus, Φ depends both on the displacement and electric potential jumps across Γ and acts as a potential jointly for the mechanical tractions and the surface-charge density.

It bears emphasis that the ferroelectric cohesive law ((6a) and (6b)) allows for an arbitrary coupling of the mechanical and electrical fields. It should also be carefully noted that the ferroelectric cohesive law describes the physics of mechanical or electrical decohesion and does not presume a particular form of the constitutive law governing the behavior in the bulk. In particular, it is possible to apply the ferroelectric cohesive law ((6a) and (6b)) in conjunction with Landau–Ginzburg–Devonshire models of domain switching (cf., e.g., Ref. [34]). The detailed boundary conditions on the crack faces are thought to greatly affect the fracture behavior of electrically driven crack growth [35]. For instance, partial discharge or charge separation effects have been suggested to play an important role in the vicinity of the crack tip [36]. The ferroelectric cohesive law ((6a) and (6b)) provides a useful framework for modeling those phenomena as well. Moreover, the ferroelectric cohesive law encodes the physics of decohesion, and thus can be tailored to represent any of the localized mechanisms of ferroelectric fatigue.

2.1. Ferroelectric fatigue cohesive law

As noted by Nguyen et al. [25], reversible cohesive laws do not predict crack advance under cyclic loading and,

therefore, are insufficient for modeling fatigue. Instead, for a cohesive law to predict fatigue it must be irreversible and account for loading–unloading hysteresis. Loading–unloading irreversibility may be built into a cohesive law by means of the internal variable formalism (cf., e.g., Ref. [30]). The loading–unloading hysteresis model developed subsequently extends that of Nguyen et al. [25] to ferroelectric fatigue.

The modeling process is greatly simplified by the assumption that the cohesive potential Φ depends on the displacement and electric-potential jumps only through the effective electro-mechanical jump [37,30]

$$\delta = \sqrt{\delta_n^2 + \beta^2 \delta_s^2 + \gamma^2 \psi^2}, \tag{7}$$

i.e., by assuming

$$\Phi = \Phi(\delta, \mathbf{q}), \tag{8}$$

where we write

$$\boldsymbol{\delta} = \llbracket \mathbf{u} \rrbracket, \tag{9a}$$

$$\delta_n = \boldsymbol{\delta} \cdot \mathbf{n}, \tag{9b}$$

$$\boldsymbol{\delta}_n = \delta_n \mathbf{n}, \tag{9c}$$

$$\delta_s = \sqrt{\boldsymbol{\delta}^2 - \delta_n^2}, \tag{9d}$$

$$\boldsymbol{\delta}_s = \boldsymbol{\delta} - \boldsymbol{\delta}_n = \delta_s \mathbf{s}, \tag{9e}$$

$$\psi = \llbracket \phi \rrbracket, \tag{9f}$$

for the normal and tangential components of the opening displacement and the electric potential jump, respectively. The parameters β and γ assign different relative weights to normal and tangential opening displacements, thus differentiating between mode I and modes II and III of fracture; and to opening displacements and the electric potential jump, thus differentiating between mechanical and electrical fatigue.

An effective electro-mechanical flux may also be defined as

$$\sigma = \frac{\partial \Phi}{\partial \delta}. \tag{10}$$

Using the chain rule, (6a) and (6b) evaluate to

$$\boldsymbol{\tau} = \frac{\sigma}{\delta} [(1 - \beta^2) \boldsymbol{\delta}_n \mathbf{n} + \beta^2 \boldsymbol{\delta}] = \frac{\sigma}{\delta} (\delta_n \mathbf{n} + \beta^2 \delta_s \mathbf{s}) = \tau_n \mathbf{n} + \tau_s \mathbf{s}, \tag{11a}$$

$$-\omega_f = \frac{\sigma}{\delta} \gamma^2 \psi. \tag{11b}$$

We also note that

$$\sigma = \sqrt{\tau_n^2 + \tau_s^2 / \beta^2 + \omega_f^2 / \gamma^2}. \tag{12}$$

Next we further specialize the preceding framework along the lines of the cohesive model of fatigue crack nucleation and growth proposed by Serebrinsky and Ortiz [26] and by Nguyen et al. [25]. The essential feature to include in the model is loading–unloading hysteresis. The specific scheme chosen to build hysteresis into the model is illustrated in Fig. 1. Monotonic loading is characterized by a cohesive

initial-boundary-value problem for the polarization \mathbf{P} , the electro-static potential ϕ and the mechanical displacement \mathbf{u} .

3. Experimental validation

In order to make contact with experiment and assess the validity of the model, we consider a simple test configuration consisting of an infinite slab of thickness u acted upon by an oscillatory voltage differential $\Delta\phi_T$ across the slab and otherwise stress free. The mid plane of the slab is a weak interface governed by the cohesive model proposed in Section 2. In order to simplify the analysis, we assume all fields to be uniform outside the cohesive interface. This assumption has the effect of reducing the initial-boundary-value problem (22)–(24) to a simple set of ordinary differential equations in time. Taking the cohesive plane to coincide with the (x_1, x_2) -coordinate plane and assuming uniaxial strain conditions, $\epsilon_{11} = \epsilon_{22} = \epsilon_{12} = \epsilon_{13} = \epsilon_{23} = 0$, the various bulk energy densities reduce to

$$W_1(P_3) = \frac{a_1}{2}P_3^2 + \frac{a_2}{4}P_3^4 + \frac{a_3}{6}P_3^6, \quad (26a)$$

$$W_2(\epsilon_{33}) = \frac{c_1}{2}\epsilon_{33}^2, \quad (26b)$$

$$W_3(P_3, \epsilon_{33}) = -\frac{b_1}{2}\epsilon_{33}P_3^2, \quad (26c)$$

where a_1 , a_2 , a_3 , c_1 and b_1 are empirical constants. The transverse stress follows from these expressions as:

$$\sigma_{33} = c_1\epsilon_{33} - \frac{b_1}{2}P_3^2. \quad (27)$$

The stress-free condition $\sigma_{33} = 0$ then gives the transverse strain as

$$\epsilon_{33} = \frac{b_1}{2c_1}P_3^2. \quad (28)$$

Using this identity, (22) reduces to

$$\mu\dot{P}_3 + a_1P_3 + \left(a_2 - \frac{b_1^2}{2c_1}\right)P_3^3 + a_3P_3^5 = E_3. \quad (29)$$

From Eq. (23) and the boundary conditions, at the cohesive interface, we additionally have

$$-\omega_f = P_3 + \epsilon_0 E_3, \quad (30a)$$

$$\psi = E_3 u + \Delta\phi_T. \quad (30b)$$

Finally, the coupled electro-mechanical cohesive law specializes to

$$\dot{\sigma} = f_1(\dot{\delta}, K_1, \sigma/\delta)\dot{\delta}, \quad (31a)$$

$$\dot{K}_1 = f_2(\dot{\delta}, K_1, \sigma/\delta)\dot{\delta}, \quad (31b)$$

$$\gamma^2 \left(\frac{P_3 u - \epsilon_0 \Delta\phi_T}{\sigma u \gamma^2 - \epsilon_0 \delta} \right)^2 = 1, \quad (31c)$$

where

$$\delta = \gamma|\psi|, \quad (32a)$$

$$\sigma = \frac{1}{\gamma}|\omega|. \quad (32b)$$

In calculations, we enforce the constraint (31c) in differential form and solve the resulting system of ordinary differential equations in time by means of a fourth-order Runge–Kutta algorithm. In implementing this algorithm, care must be exercised in order to resolve ambiguities in the selection of branches and the handling of singularities in the response functions as δ tends to zero. In the sequel, $P = P_3$ and similarly for other vector quantities, and $\epsilon = \epsilon_{33}$.

For sufficiently high applied field, the calculated response exhibits characteristic hysteresis loops in a P – E plot, and butterfly loops in a ϵ – E plot. These loops are shown in Fig. 2 for several values of the peak electric field E_p and the applied-field frequency f . Several aspects of the bulk response are noteworthy. Thus, below a minimum value of E_p no switching occurs, and the sign of the polarization

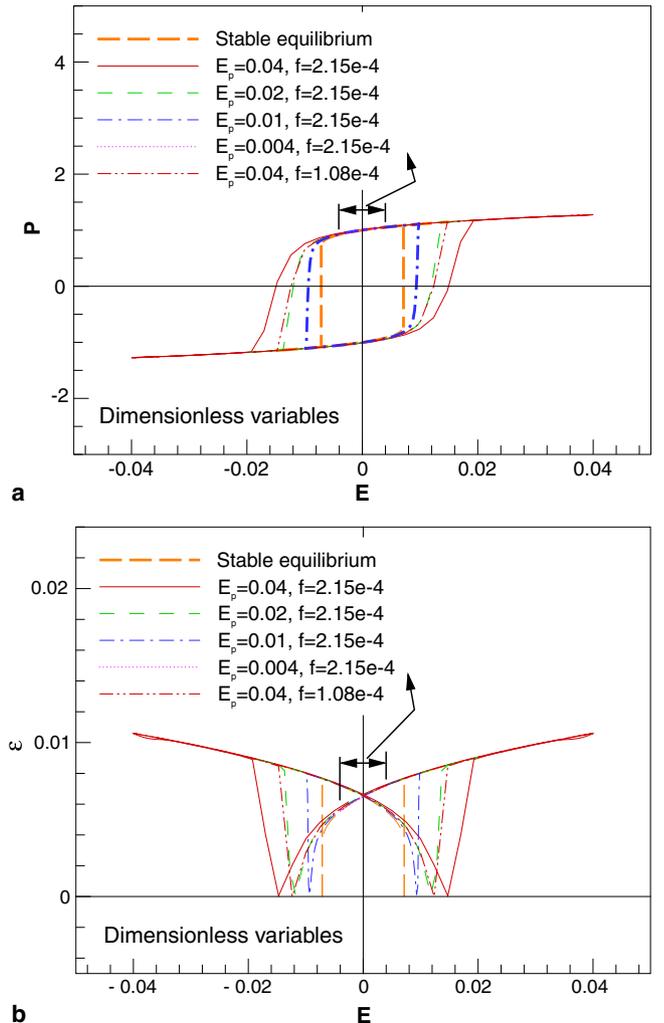


Fig. 2. Bulk response for different field amplitudes and frequencies. (a) P – E hysteresis loops. (b) ϵ – E butterfly loops.

remains constant in time. The rate-dependency introduced by the time-dependent Ginzburg–Landau equation is evident in Fig. 2. In the limit of $\mu = 0$ or, equivalently, of a small frequency of the applied field, this rate dependency is removed and an equilibrium loop is obtained. For non-zero μ , the bulk response depends on the applied-field frequency.

The predictions of the model in the example just described can now be compared with experimental data by way of validation. In order to facilitate comparisons for different materials, we introduce the following normalization constants for the variables: polarization, P_{ref} ; stress, elastic constants, c_{ref} ; permittivity, $\epsilon_{0\text{ref}} = P_{\text{ref}}^2/c_{\text{ref}}$; electric field, $E_{\text{ref}} = P_{\text{ref}}/\epsilon_{0\text{ref}}$; thickness, displacements, $L_{\text{ref}} = \delta_a$; potential, $\phi_{\text{ref}} = E_{\text{ref}}L_{\text{ref}}$; time $t_{\text{ref}} = \mu P_{\text{ref}}^2/c_{\text{ref}}$; energy density coefficients, $b_{\text{ref}} = a_{1\text{ref}} = c_{\text{ref}}/P_{\text{ref}}$, $a_{2\text{ref}} = a_{1\text{ref}}/P_{\text{ref}}^2$, $a_{3\text{ref}} = a_{1\text{ref}}/P_{\text{ref}}^4$; coefficient $\gamma_{\text{ref}} = P_{\text{ref}}/c_{\text{ref}}$; cohesive stiffness $K_{\text{ref}} = c_{\text{ref}}/L_{\text{ref}}$. Dimensionless variables are used in the calculations, and the parameters and material constants are summarized in Table 1. We have selected material constants for the bulk model readily available from the literature [38], which correspond to BaTiO₃. The initial conditions are $P_{30} = 1$, $K_{10} = 2311$, and δ_0 and σ_0 are chosen to give a ratio $K_{10}/K_{u0} = 0.99$.

Typical experimental data are given in the form

$$P_x = P_x(E_p^*, N, u), \quad (33)$$

where $E_p^* = -\Delta\phi_p/u$ is the peak *nominal* electric field, which should be carefully differentiated from the peak *actual* field E_p in the material, N is the cycle number, and P_x is a measure of the polarization state of the material, typically the remanent polarization P_r but also sometimes P_{sw} , the switching polarization. From these data, a relation $E_f^*(N, u)$, or the inverse relation $N_f(E_p^*, u)$, can be derived by solving the equation

$$\frac{P_x(E_p^*, N, u)}{P_x(E_p^*, 1, u)} = 1 - C, \quad (34)$$

where C is the loss of polarization due to fatigue. We shall take these fatigue maps as the basis for the validation of the model.

As expected, the predicted fatigue behavior depends critically on whether switching occurs. In the presence of switching, the interface degrades upon cycling and the peak value of δ increases monotonically with the number of cycles, Fig. 3. The increase in δ is initially slow and accelerates markedly in the last stages of the fatigue life of the material. The rate of damage accumulation increases with the amplitude of the nominal field. Correspondingly, as the interface degrades the switching capability and the actuation strain are impaired.

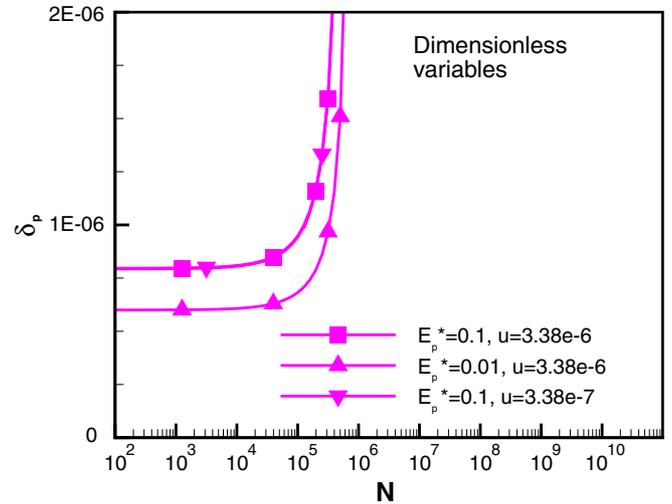


Fig. 3. Evolution with number of cycles of the peak effective opening displacement δ_p .

The calculated fatigue map, for small frequencies for which rate-dependency is negligible, is shown in Fig. 4(a). It is immediately apparent from this map that for each thickness u there is a threshold amplitude $E_{f,\text{th}}^*$ below which fatigue does not occur. Specifically, fatigue occurs if the nominal field E_p^* is sufficiently strong to cause switching, and fatigue does not occur otherwise. Thus, the fatigue threshold $E_{f,\text{th}}^*$ coincides with the nominal coercive field E_{cv}^* for the virgin material. For nominal fields above the fatigue threshold $E_{f,\text{th}}^*$, the model predicts the fatigue life N_f to shorten with increasing nominal field amplitude E_p^* . This dependency is well approximated by the power law

$$N_f = AE_p^{*n}, \quad (35)$$

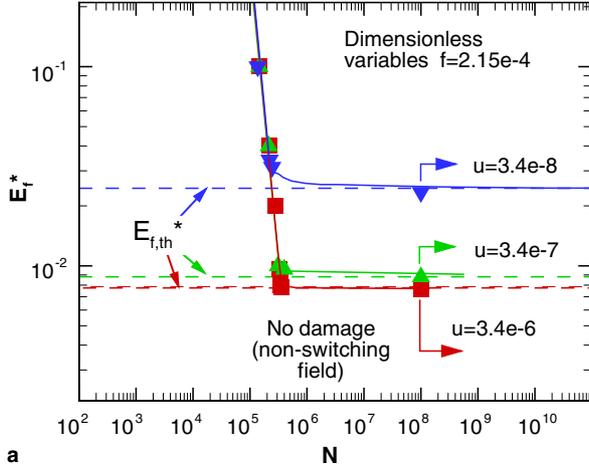
with exponent $-n \approx 2.8$.

For bulk-like samples and high frequencies, the fatigue life has a steeper dependency on the applied field, as shown in Fig. 5(a). It is readily verified that the transition from non-switching to switching bulk behavior, as the applied field increases, appears smoother as the field frequency increases, thus leading to the calculated fatigue behavior. In addition, $E_{f,\text{th}}^*$ can be greatly in excess of E_{cv}^* , as Fig. 5(b) shows. It also indicates that the low frequency limit is attained for the lowest frequency used.

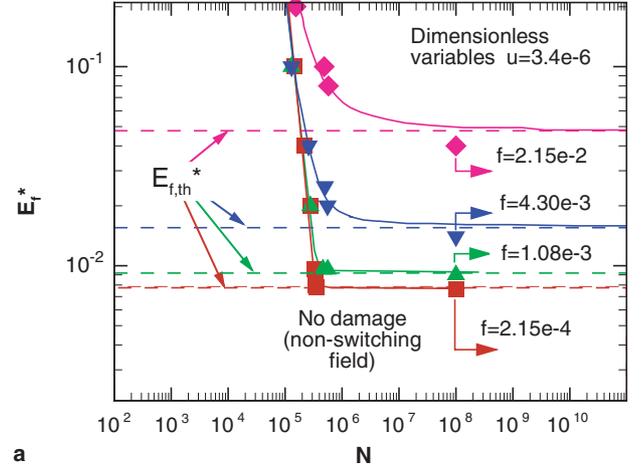
Corresponding experimental data for smooth tetragonal PZT samples [39,1,40] are shown in Fig. 4(b) for purposes of comparison. Evidently, the predicted existence of a fatigue threshold and the reduction in fatigue life with nominal field amplitude are consistent with the data. The nominal fatigue threshold $E_{f,\text{th}}^*$ is indeed observed to be greater than or equal to the nominal coercive field E_{cv}^* , as obtained from the respective hysteresis loops. Besides these experiments where no microcracking was observed or analyzed, the relation between switching and fatigue has also been established on indented and pre-cracked samples by the experiments of Zhu et al. [22]. Thus, switching appears as a necessary (though not suffi-

Table 1
Values of the model parameters

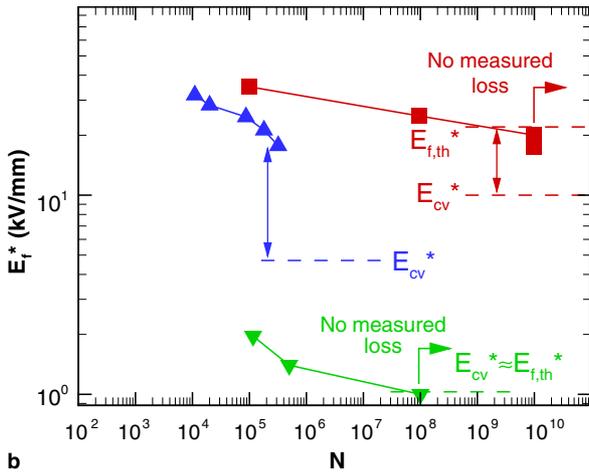
a_1	a_2	a_3	b_1	c_1	ϵ_0	γ	μ
-0.007	-0.009	0.0261	1.4282	185	0.131	800	1



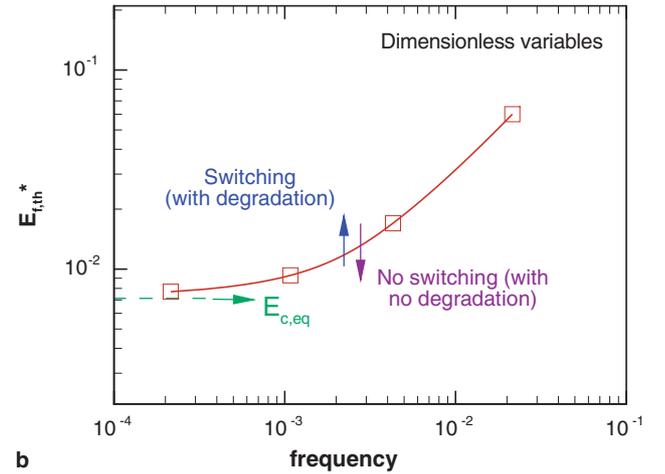
a



a



b



b

Fig. 4. Fatigue life as a function of nominal field and thickness. (a) Calculated. In all cases, $E_{f,th}^* = E_{cv}^*$. (b) Experimental (all in PZT), loss = 10%. ■: Grossmann et al. [39], ▼: Nuffer et al. [1], ▲: Mihara et al. [40]. In all cases, $E_{f,th}^* \geq E_{cv}^*$.

cient) condition for electrical fatigue, regardless of the fatigue mechanism operating. It should be noted that there are other methods, alternative to electric fatigue, to suppress polarization (e.g., [41]). The experimental data shown in Fig. 4(b) is also suggestive of power-law behavior (see Eq. (35)) with exponents in the range $-n \approx 0.16$ – 0.24 . The difference with the predicted value is due to the choice bulk behavior, which has no dependence of the actual coercive field on actual field amplitude for $E_p > E_c$, and not to the cohesive model. The dependence of the fatigue threshold on the applied-field frequency, with high frequencies delaying the onset of fatigue, was established by Grossmann et al. [39].

It is worth mentioning that some alternatives were developed to suppress polarization fatigue, including using conducting oxide electrodes [42]. Such devices can be modeled by an interface with a large value of δ_a , which would lead to a minor fatigue effect. In the limit of $\delta_a \rightarrow \infty$, fatigue would be completely absent.

In order to exhibit the size effect predicted by the model, Fig. 6(a) plots specimen thickness against the minimum

Fig. 5. Effect of applied field frequency on fatigue properties. (a) Fatigue life as a function of applied field frequency. (b) Effect of frequency on nominal threshold field.

value of E_p^* for which switching occurs, and the maximum value of E_p^* for which switching does not occur. Insight into this dependence may be derived by analyzing the equilibrium case of $\dot{P} = 0$ in Eq. (22). In this case, the governing equations reduce to

$$E = a_1 P + \left(a_2 - \frac{b_1^2}{2c_1} \right) P^3 + a_3 P^5, \quad (36a)$$

$$P = E(K_u \gamma^2 u - \epsilon_0) + K_u \gamma^2 \Delta \phi_T. \quad (36b)$$

For a virgin material, i.e., a material with high K_u , it is readily shown that the nominal coercive field E_{cv}^* required for switching is of the form

$$E_{cv}^*(u) = \Delta \phi_c(u)/u \sim E_{ceq} + (P_{ceq} + \epsilon_0 E_{ceq}) / (K_u \gamma^2 u). \quad (37)$$

where $E_{ceq} = 0.007129$. This relation is plotted in Fig. 6(a) along with the exact calculated values of $E_{cv}^*(u)$ from Eq. (36a). It is noteworthy that E_{cv}^* decreases monotonically with size u to a limiting value E_{ceq} characteristic of large components. Conversely, E_{cv}^* increases as u^{-1} for a small component thickness. This prediction of the model is born out by the experimental data, Fig. 6, which is

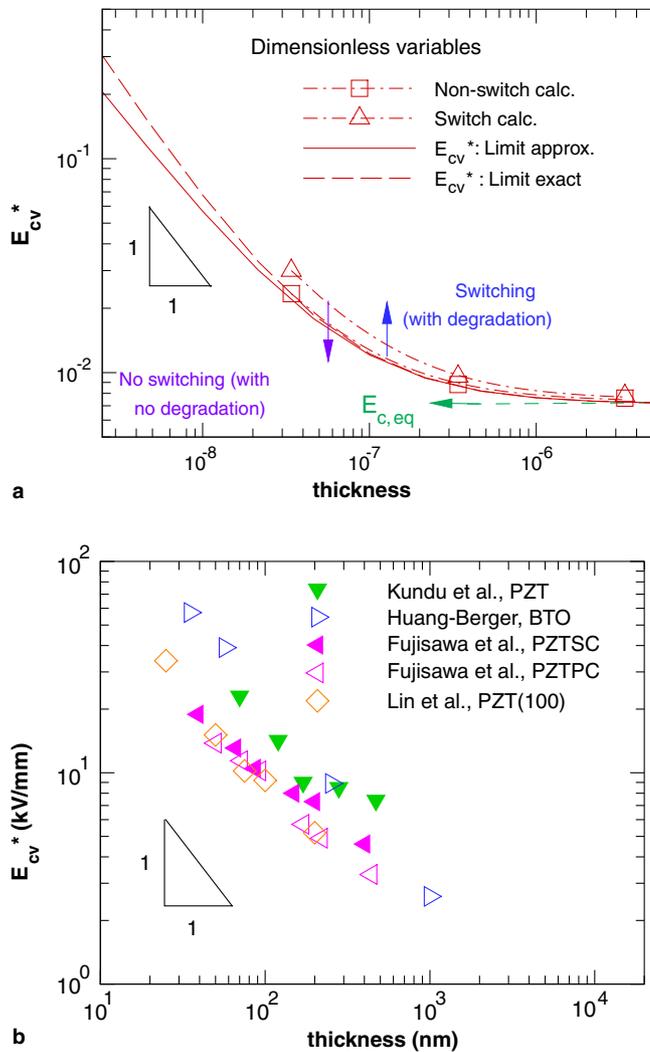


Fig. 6. Nominal electric field required for switching. (a) Calculated. Determined from Fig. 4(a): \triangle : lowest values of E_p^* with switching behavior, \square : highest values of E_p^* with non-switching behavior. Determined from models with $K_1 = K_u = \text{const.}$: - : approximate model, - - : exact model. (b) Experimental. \blacktriangleleft , \triangleleft : Fujisawa et al. [43], \blacktriangledown : Kundu and Lee [44], \triangleright : Huang and Berger [45], \diamond : Lin et al. [46].

consistent with the $E_{cv}^* \sim u^{-1}$ scaling behavior. By contrast, the size effect above the nominal fatigue threshold $E_{f,th}^*$ is negligible, Fig. 4(a). This lack of sensitivity to the component size is also visible in the evolution of δ_p , Fig. 3. Thus, the main effect of component size is on the ability of the material to switch and, by extension, on the nominal fatigue threshold $E_{f,th}^*$. It should be noted that alternative explanations for the size effect have been proposed, including depolarization fields [47,48], epitaxial stress effects [49,45] and variants of a conductive layer next to the ferroelectric–electrode interface [50,51].

4. Summary and concluding remarks

We have presented a model of electro-mechanical ferroelectric fatigue based on the postulate of a ferroelectric cohesive law that: couples mechanical displacement and

electric-potential discontinuity to mechanical tractions and surface-charge density; and exhibits a monotonic envelope and loading–unloading hysteresis. In conjunction with a constitutive model accounting for domain switching, the electro-mechanical cohesive fatigue law is able to induce ferroelectric fatigue by the following mechanism: as degradation proceeds, the surface of electro-mechanical discontinuity absorbs an increasingly large amount of the displacement and/or electric potential difference, thereby unloading the bulk and hindering switching. We identify the end of the fatigue life with the time at which the material loses its ability to sustain loading/applied field cycles of a certain constant amplitude. We have compared selected predictions of the model with experimental data for a simple test configuration consisting of an infinite slab acted upon by an oscillatory voltage differential across the slab and otherwise stress free. The model captures salient features of the experimental record including: the existence of a threshold nominal field for the onset of fatigue; the dependence of the threshold on the applied-field frequency; the dependence of fatigue life on the amplitude of the nominal field; and the dependence of the coercive field, and thus of the fatigue threshold, on the size of the component, or size effect. Our results seem to indicate that planar-like regions affected by cycling may lead to the observed fatigue in tetragonal PZT.

The ability of the model to predict the observed size effect stems directly from the fact that cohesive laws introduce a characteristic or intrinsic material length scale. Hence, in the present model the size effect is a material property and a direct consequence of material behavior. This is in contrast to other explanations of the size effect found in the literature (e.g., [50,51,49]), where the length scale is often structural.

In closing, we emphasize that the primary focus of this study has been to investigate qualitative trends and no systematic attempt has been made to optimize fit to the experimental data. It is conceivable that good quantitative agreement with the data could be obtained by some extensions and careful calibration of the model, including full finite element calculations; the use of more accurate bulk constitutive relations; and a systematic identification of aspects of the cohesive law such as the precise shape of the monotonic envelope and of the loading–unloading law.

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