The Anisotropy Induced by a Magnetostriction in Exchange-Biased Two-Layer Films

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The exchange bias at ferromagnetic (FM)/antiferromagnetic (AF) interfaces strongly depends upon the state of antiferromagnetic layer, which is sensitive to mechanical stresses due to its strong magnetoelastic coupling. In a given paper, we consider magnetoelastic effects, which arise at FM/AF interface due to misfit of lattices and magnetic ordering. We show how magnetostriction affects mutual orientation of the AF and FM vectors as well as magnetic easy-axis direction in thin AF layer. The results obtained can be used for tailoring of exchange-biased systems.
ориентацию $\Phi$- и $\Phi M$-векторов и на направление лёгкой оси в тонком $\Phi F$-слое. Полученные результаты могут быть использованы при создании систем с обменным подмагничиванием.

**Key words:** magnetic anisotropy, exchange bias, magnetoelasticity, magnetic properties of interfaces, antiferromagnet.

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1. INTRODUCTION

Antiferromagnetic (AF) materials are widely used in spintronic devices as auxiliary elements for pinning of ferromagnetic (FM) magnetization through the effect of exchange bias (see, *e.g.* [1]). The possibility to control the state of coupled AF/FM bilayers requires investigation of the magnetic mechanisms that could be responsible for bias effect. Many researchers [2—7] emphasize the important role of the AF domain structure in the establishing of the exchange bias. The problem of AF domains is intimately related with magnetoelastic coupling [8] and can strongly depend upon the mechanical stress that appears at the FM/AF interface due to the lattice misfit. Magnetostriction can also provide additional coupling between FM and AF layers and affect orientation of AF moments in the near-surface region [9, 10]. The widely studied epitaxial films consisting of FM and nonmagnetic materials [11—14] show strong correlation between magnetoelastic coupling and magnetic properties. Analogous and even more striking phenomena could be expected in the systems, which combine FM with AF that possesses large magnetostriction.

In the present paper, we show that magnetostriction of AF produces uniaxial anisotropy in the plane of the adjacent FM layer and, thus, causes strong surface magnetic anisotropy in AF itself.

2. UNIAXIAL ANISOTROPY OF FERROMAGNET

Epitaxial ferromagnetic film deposited on top of AF inherits the crystallographic structure of the substrate. If the substrate has a certain anisotropy induced by magnetoelastic strains, this anisotropy in atomic arrangement will be reproduced by the FM layer. Thus, additional contribution to the magnetic energy of film should be proportional to magnetoelastic coupling in both FM and AF materials. Phenomenological expression for such a type of uniaxial in-plane anisotropy can be deduced from the magnetoelastic energy of FM, which for a cubic-symmetry crystal is as follows:

\[
\mathcal{E}_{me} = b_1^f \left( u_{xx} \alpha_x^2 + u_{yy} \alpha_y^2 + u_{zz} \alpha_z^2 \right) + 2b_2^f \left( \alpha_x \alpha_y u_{xy} + \alpha_y \alpha_z u_{yz} + \alpha_z \alpha_x u_{zx} \right). \tag{1}
\]
Here, \( u_{ik} \) are strain tensor components, which we calculate with respect to the bulk nonmagnetic reference state, \( b_{k} \) are magnetoelastic-coupling coefficients. Magnetisation vector \( \mathbf{M}_F \) of FM is described by the direction cosines \( \alpha_k, k = x, y, z \). In the relaxed state of FM/AF system, an equilibrium strain \( u_{ik} \) includes deformations produced by lattice mismatch and spontaneous strain \( \bar{u}_{\text{mag}} \) induced by magnetic ordering in the AF substrate. For a symmetric (001) surface, the misfit-induced strains are isotropic and can influence only out-of-plane anisotropy of FM. In contrast, magnetostrictive contribution, though small as compared with the misfit strain, has nontrivial shear components, \( u_{xx}^{AF} - u_{yy}^{AF} \) and/or \( u_{xy}^{AF} \) which can remove degeneracy between different in-plane directions. Thus, uniaxial contribution into magneto-crystalline energy of FM film takes a form

\[
f_a^{xx} = \frac{1}{2} K_{1ua}^{F} \rho_j (\alpha_x^2 - \alpha_y^2) + K_{2ua}^{F} \rho_j \alpha_x \alpha_y
\]

with anisotropy constants

\[
K_{1ua}^{F} = b_1^{F} (u_{xx}^{AF} - u_{yy}^{AF}), \quad K_{2ua}^{F} = 2b_2^{F} u_{xy}^{AF}.
\]

Variable \( \rho_j = \pm 1 \) distinguishes between the different domains of AF.

A preferable direction of FM magnetisation \( \mathbf{M}_F \), which depends upon the sign of the coefficients \( K_{ua}^{F} \), is defined by correlation between selfstriction of the FM and external striction imposed by the AF. If, for example, magnetostriction of FM in the direction of magnetisation is positive (elongation, \( b^F < 0 \)), \( \mathbf{M}_F \) will tend to align in the direction of maximal elongation of the AF, i.e., for positive \( u^{AF} \) value (elongation), \( K_{ua}^{F} \) is negative, as can be easily checked from equation (3).

Magnetostriction-induced uniaxial anisotropy (3) competes with the anisotropy arising from the FM/AF exchange in a thin near-surface region of thickness \( \xi \). For a compensated AF surface, this contribution depends upon the exchange integral between the atoms of FM and AF, \( J_{F-AF} \), and susceptibility of AF, \( \chi_{AF} = 1/J_{AF} \) (Koon’s model, [15]):

\[
f_{\text{exch}} = -\frac{1}{2} \chi_{AF} J_{F-AF}^2 [\mathbf{M}_F \times \mathbf{L}_S]^2.
\]

The AF vector \( \mathbf{L}_S \) describes orientation of spins at the surface of the AF substrate (which in principle can differ from that in the bulk, as will be shown later).

To elucidate the effect of both contributions, let us consider a simple case when one of the in-plane easy axes (say, \( x \)) of FM coincides with in-plane \( \mathbf{L}_S \) direction and \( u_{xy}^{AF} = 0 \). For the in-plane FM ordering (\( \alpha_z = 0 \)), we set \( \alpha_x = \cos \psi, \alpha_y = \sin \psi \). Thus, the effective energy is
\[ f_{\text{eff}}^V = \frac{1}{4} K_4 \sin^2 2\psi + \frac{1}{2} [K_{uu}\rho_j + \frac{\zeta}{2t_F} \chi_{AF}J_{AF}^2] \cos 2\psi. \] 

(5)

Constant \( K_4 > 0 \) is magnetocrystalline constant, and we suppose the FM film to be homogeneously ordered throughout the thickness \( t_F \).

Equilibrium value \( \psi = \psi_{eq} \) minimizes effective energy (5), so, it satisfies the relations

\[ \sin 2\psi_{eq}\{K_4 \cos 2\psi_{eq} - [K_{uu}\rho_j + \frac{\zeta}{2t_F} \chi_{AF}J_{AF}^2]\} = 0. \] 

(6)

\[ K_4 \cos 4\psi_{eq} - [2K_{uu}\rho_j + \frac{\zeta}{t_F} \chi_{AF}J_{AF}^2] \cos 2\psi_{eq} > 0. \] 

(7)

In the absence of FM/AF interaction, FM has two equivalent easy directions in (001) plane, \( \psi_1^{(0)} = 0 \) and \( \psi_2^{(0)} = \pi/2 \). Antiferromagnetic substrate removes this degeneracy. If exchange coupling is not too large, \( \frac{\zeta}{t_F} \chi_{AF}J_{AF}^2 \leq K_4 t_F \), both solutions \( \psi_{1,2}^{eq} \) satisfy equations (6), but have different energies, the difference being

\[ f_{\text{eff}}^V(\psi_{1,2}^{eq}) - f_{\text{eff}}^V(\psi_{3,4}^{eq}) = K_{uu}\rho_j + \frac{\zeta}{2t_F} \chi_{AF}J_{AF}^2. \] 

(8)

It can be easily seen from (8) that the FM/AF exchange coupling makes favourable the solution with \( M_F \perp L_S \) (\( \psi_2^{eq} = \pi/2 \)) for any sign of the exchange constant \( J_{AF} \). In turn, magnetostriction-induced anisotropy \( K_{uu} \) can oppose this tendency and can make preferable in-paralle of orientation of \( M_F \) and \( L_S \) (\( \psi_1^{eq} = 0 \)). It should be stressed that these two mechanisms have different origin and the system can switch from one easy-axis to another with variation of FM thickness. Exchange mechanism ties together mutual orientation of FM magnetisation and AF spins in the near-surface layer. This mechanism is important for very thin films, where factor \( \frac{\zeta}{t_F} \) is not vanishingly small. Magnetostriction-related mechanism is a long-range one; it depends upon orientation of AF moments in the bulk, which can be different from \( L_S \).

Moreover, in some AFs widely used in FM/AF systems (e.g., NiO, CoO, LaFeO$_3$, KCoF$_3$), magnetostriction originates from the strong spatial dependence of the exchange integral and is insensitive to exact orientation of AF spins. In this very important case, uniaxial anisotropy of FM is defined mainly by the domain structure of AF.

The role of magnetostriction-induced mechanism can be illustrated by some experimental examples. Simultaneous observation of the FM and AF spins in Co/LaFeO$_3$ [16] and Co/NiO [17] systems revealed that FM magnetisation is aligned parallel or antiparallel to the in-plane projection of the AF axis in contrast to the usually observed perpendicular
coupling consistent with the Koon’s model [15]. Uniaxial anisotropy is also detected after deposition of Fe on top of KCoF₃ [18, 19]. All these AFs are known to have rather large magnetostriction of the exchange nature (see Table 1).

Using the values of magnetoelastic constants for ferromagnets (Table 2), one can calculate from equation (3) the expected value of uniaxial anisotropy in different FM/AF combinations (see Table 3).

As can be seen from Table 3, uniaxial anisotropy in the Fe film constitutes only \( \approx 10\% \) from the ‘pure’ magnetic anisotropy. Nevertheless, this value will last to choose preferable axis of magnetisation as was clearly observed in the experiments [18, 19].

**TABLE 1.** Magnetostriction (spontaneous deformations) of typical AFs calculated from the experimentally observed lattice constants above and below Néel temperature.

<table>
<thead>
<tr>
<th>AF</th>
<th>Magnetostriction</th>
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<tbody>
<tr>
<td>NiO</td>
<td>(-2.6 \cdot 10^{-3}) [20]</td>
</tr>
<tr>
<td>LaFeO₃</td>
<td>(-4.76 \cdot 10^{-4}) [21]</td>
</tr>
<tr>
<td>KCoF₃</td>
<td>(-2.0 \cdot 10^{-3}) [22, 23]</td>
</tr>
<tr>
<td>CoO</td>
<td>(-2 \cdot 10^{-2}) [24]</td>
</tr>
</tbody>
</table>

**TABLE 2.** Magnetoelastic coupling coefficients for FMs [11] (in [erg/cm³]).

<table>
<thead>
<tr>
<th></th>
<th>Co, f.c.c.</th>
<th>Fe, b.c.c.</th>
</tr>
</thead>
<tbody>
<tr>
<td>(b₁)</td>
<td>(-9.2 \cdot 10^7)</td>
<td>(-3.43 \cdot 10^7)</td>
</tr>
<tr>
<td>(b₂)</td>
<td>(7.7 \cdot 10^7)</td>
<td>(7.83 \cdot 10^7)</td>
</tr>
</tbody>
</table>

**TABLE 3.** Magnetic anisotropy of systems [erg/cm³]. \(K₄\) (2nd column) is the 4th order magnetocrystalline anisotropy observed in the bulk Fe and Co crystals. Theoretical values of \(K_{ua}\) (third column) are calculated from Eq. (3). Experimental values of \(K_{ua}\) (the last column) are extracted from measurement of hysteresis loops (for Co) and ferromagnetic resonance (for Fe).

<table>
<thead>
<tr>
<th>FM/AF system</th>
<th>(K₄)</th>
<th>(K_{ua})</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Theory</td>
<td>Experiment</td>
</tr>
<tr>
<td>Co/NiO</td>
<td>(-2.3 \cdot 10^5) [26]</td>
<td>(2.0 \cdot 10^5)</td>
</tr>
<tr>
<td>Co/LaFeO₃</td>
<td>(0.37 \cdot 10^5)</td>
<td>(1.4 \cdot 10^5)</td>
</tr>
<tr>
<td>Co/CoO</td>
<td>(6.0 \cdot 10^6)</td>
<td>(1.2 \cdot 10^6)</td>
</tr>
<tr>
<td>Fe/NiO</td>
<td>(8.5 \cdot 10^5) [12, 19]</td>
<td>(0.9 \cdot 10^5)</td>
</tr>
<tr>
<td>Fe/KCoF₃</td>
<td>(0.7 \cdot 10^5)</td>
<td>(0.8 \cdot 10^5) [19]</td>
</tr>
</tbody>
</table>
More pronounced effect is expected in Co films, which have rather high magnetostriction and small bulk magnetic anisotropy. Predicted value of the uniaxial anisotropy is of the same order as $K_4$ or even one order of magnitude larger, than in the case of Co/CoO. It should be noted that in calculation we started from the bulk values of magnetoelastic coefficients for Fe and Co. In the case of ultrathin Co films, these values need to be ascertained because of the large potential misfit between FM and AF lattices (nearly 10%). Depending on the growth mode, this mismatch can either relax through the formation of dislocations, or produce strong internal stresses in the Co film, which, in turn, can give rise to a crucial change of the value and even the sign of magnetoelastic coefficient (see, e.g., Refs. [13, 25]).

3. SURFACE ANISOTROPY OF ANTIFERROMAGNET

It is widely recognised that lattice misfit strongly influences the magnetic and magnetoelastic properties of the film (see, e.g. [29]). On the other hand, epitaxial misfit may equally induce large stress in the substrate (this phenomenon is used to measure stress in the film [11]). In the case when the substrate is rather thick, stress exerted by the film relaxes over a small distance $\Sigma^{AF}$ in the near-surface layer of AF. For AFs with large magnetoelastic coupling, this surface stress can produce an additional magnetic anisotropy, which we will call a surface anisotropy.

Phenomenological description of this effect is based on the analysis of the Helmholtz free energy potential $G$, which includes elastic $f_e$ and magnetoelastic $f_{me}^{AF}$ energy of AF using antiferromagnetic vector $L$ and components of stress tensor $\sigma$ as the internal parameters:

$$G = \int (f_{me}^{AF} + f_e) dV.$$  \hspace{1cm} (9)

In the simplest case of a cubic crystal, the elastic energy density $f_e$ takes a form

$$f_e = \frac{s_{11}}{2} \left( \sigma_{xx}^2 + \sigma_{yy}^2 + \sigma_{zz}^2 \right) + s_{12} \left( \sigma_{xx} \sigma_{yy} + \sigma_{yy} \sigma_{zz} + \sigma_{zz} \sigma_{xx} \right) + 2s_{44} \left( \sigma_{xy}^2 + \sigma_{yz}^2 + \sigma_{zx}^2 \right),$$  \hspace{1cm} (10)

where we turned from strains to stresses using the Hooke’s law. Compliances $s_{ik}$ are expressed through the elastic moduli $c_{ik}$ in a usual way:

$$s_{11} = \frac{c_{11} + c_{12}}{(c_{11} - c_{12})(c_{11} + 2c_{12})}, \hspace{0.5cm} s_{12} = -\frac{c_{12}}{(c_{11} - c_{12})(c_{11} + 2c_{12})}, \hspace{0.5cm} s_{44} = \frac{1}{c_{44}}.$$  \hspace{1cm} (11)
Density of magnetoelastic energy $f_{me}$ can be written as

$$f_{me}^{AF} = f_{exch}^{AF} + \frac{b_{11}^{AF} + b_{12}^{AF}}{c_{11} - c_{12}} \left( \sigma_{xx} L_x^2 + \sigma_{yy} L_y^2 + \sigma_{zz} L_z^2 \right) +$$

$$+ 2 \frac{b_{44}^{AF}}{c_{44}} \left( L_x L_y \sigma_{xy} + L_y L_z \sigma_{yz} + L_z L_x \sigma_{zx} \right),$$

where $b_{1,2}^{AF}$ are magnetoelastic coupling coefficients of a cubic AF, and far from the Néel temperature, AF vector can be normalised, so $|L| = 1$. The first term in (12) describes a possible non-isomorphic contribution, which arises from the space dependence of the exchange interactions, described by a coefficient $B_{0}^{AF}$. It depends on the specific type of AF. For example, for a single-domain NiO, it can be expressed as

$$f_{exch}^{AF} = \frac{B_{0}^{AF}}{c_{44}} (\sigma_{xy} + \sigma_{yz} + \sigma_{zx}) L^2.$$

In the presence of the FM coverage, the AF substrate exerts a surface stress $\tau^{AF}$ opposite to the surface stress in the FM film $\tau^{F}$:

$$\tilde{\tau}^{AF} \equiv \int \sigma dz = -\tau^{F}. \quad (13)$$

$z$-axis is directed along the film normal and integration is over the AF thickness. For a (001) cubic surface, $\tau^{F}$ can be estimated from the misfit value $\epsilon_{MF}$ as follows:

$$\tau_{xx}^{F} = \tau_{yy}^{F} = \tau_{zz}^{F} = t_{F} \left( c_{11} + c_{12} - \frac{c_{12}^2}{c_{11}} \right) \epsilon_{MF}, \quad (14)$$

where $t_{F}$ is the FM film thickness. Substituting (13) into free energy (9), we obtain a contribution from the FM/AF misfit as

$$G_{eff} = \int_{S} \frac{b_{44}^{AF}}{c_{11} - c_{12}} \tau^{F} L_z^2 dS = \frac{1}{2} \int_{S} K_{S}^{AF} L_z^2 dS, \quad (15)$$

which could be associated with the surface/interface energy of AF. Effective constant

$$K_{S}^{AF} = 2b_{11}^{AF} c_{MF} \left( 1 + 2 \frac{c_{12}}{c_{11}} \right) t_{F} \quad (16)$$

is proportional to the product of magnetoelastic coupling coefficient of AF and misfit (or effective stress) in the FM layer.

The sign of $K_{S}^{AF}$ and, hence, the character of the induced surface anisotropy, is defined by the relation between the sign of AF spontaneous
striction and that of external stress. Let us suppose FM lattice constant is smaller than that of AF ($\tau^F > 0$). Then, AF surface exerts a compressive stress. According to the general Le Chatelier’s principle, AF vector at the surface will rotate in a way, which reduces the external influence. In the case of positive striction (AF spontaneously elongates in spin direction), in-plane orientation of AF spins will be preferable ($K^A_{S} > 0$). It worth to mention that the analogous, magnetoelastic, mechanism related with the rotation of AF moments in near-surface region is responsible for the shape-induced magnetic anisotropy in AF nanoparticles [30].

4. DISCUSSION

The misfit-induced surface anisotropy can produce a noticeable rotation of AF spins in the vicinity of interface region. The most pronounced effect can be expected for NiO, CoO, and LaFeO$_3$ AFs, in which the bulk AF vector makes some angle with (001) surface. Particularly, in NiO and CoO, the AF spins are ordered in (111) planes (with small deflection in the case of CoO [31–33]), in which they can be easily rotated. An easy-axis is directed along $\langle 2\bar{1}1 \rangle$ in NiO and $\langle 3\bar{1}1 \rangle$ in CoO, thus, for a cleaved (001) surface, AF moments have nonzero component perpendicular to the surface plane, as is observed for NiO crystal [34, 35].

Depositions of Fe and Co on NiO, and Fe$_3$O$_4$ and Co on CoO produce compressive surface stress in AF (see Table 4).

For NiO and CoO, the magnetoelastic constants $b^A_{F}$ are positive (as deduced from the data [24, 36]), so, as it follows from (15), (16), for all the mentioned FM/AF combinations, the preferable orientation of AF vector $L$ is in the interface (001) plane. A compromise between the strong dipole–dipole anisotropy, which tends to keep AF moments close to (111) plane, and strain-induced surface anisotropy in (001) plane is the direction $[1\bar{1}0]$. Therefore, depending on the balance be-

| TABLE 4. Bulk lattice parameters (2nd column) and calculated interatomic distances at (001) surface of f.c.c. lattice (3rd column) for different FM and AF (in [Å]). |
|---------------------------------|---------|---------|
| Fe, b.c.c.                      | 2.866 [26] | 4.053   |
| Co, f.c.c.                      | 3.544 [26] | 3.544   |
| NiO                             | 4.177    | 4.177   |
between the bulk magnetic anisotropy and the induced surface anisotropy (16), AF moments may rotate from the bulk easy direction [110] to a smaller or larger angle. The effect should be obviously stronger for Co FM because of the large misfit value.

Experimentally this phenomenon was observed in [17], where deposition of 2 nm Co film on the (001) surface of NiO resulted in the total reorientation of NiO spins to [110] direction. An observed collinear alignment of Co and NiO spins in this system arises from both misfit-induced reorientation of AF moments and magnetostriction-induced uniaxial anisotropy in the FM layer.

A similar effect was observed in the Fe3O4/CoO multilayers [3], where an influence of the surface stress is much more pronounced. In this system, all of the AF Co moments lie along [110] or [110] directions (depending on the AF domain type). This orientation does not vary with temperature, magnetic field and thickness of CoO layers.

Misfit-induced anisotropy of AF layer depends upon the internal stresses $\tau^\nu$ in the adjacent ferromagnet, which could relax in the course of field cycling. Variation of stress, in turn, affects the domain structure of AF. Therefore, magnetoelastic mechanism can explain training effect (irreversible changes in configuration of AF domains), frequently observed in bilayers with multidomain state of AF in the as-cast sample (see, e.g., Refs. [2, 37]).

In summary, we have studied the effect of magnetostriction on the properties of FM/AF coupled system. Spontaneous striction, which appears in antiferromagnet due to the AF ordering, can cause uniaxial in-plane anisotropy in the ferromagnetic film and set preferential easy axis of FM either along with or perpendicular to the orientation of AF vector. Competition between uniaxial anisotropy induced by long-range magnetostriction and short-range exchange mechanism results in different orientation of the FM easy-axis depending on the thickness of FM layer. Lattice misfit between FM and AF is a source of a magnetic surface anisotropy in AF substrate, which can cause rotation of AF moments in the near-surface region compared with their bulk orientation.

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