Equivalence of magnetoelastic, elastic, and mechanical work energies with stress-induced anisotropy

Chaitanya Mudivarthi\textsuperscript{a}, Supratik Datta\textsuperscript{b}, Jayasimha Atulasimha\textsuperscript{c}, Alison B. Flatau\textsuperscript{a,b}, Phillip G. Evans\textsuperscript{d}, and Marcelo J. Dapino\textsuperscript{d}

\textsuperscript{a}Material Science and Engineering, University of Maryland, College Park, MD-20742
\textsuperscript{b}Aerospace Engineering, University of Maryland, College Park, MD-20742
\textsuperscript{c}Department of Mathematics, North Carolina State University Raleigh, NC 27695
\textsuperscript{d}Mechanical Engineering, the Ohio State University, Columbus, OH 43210

ABSTRACT

This work investigates the equivalence of thermodynamic potentials utilizing stress-induced anisotropy energy and potentials using elastic, magnetoelastic, and mechanical work energies. The former is often used to model changes in magnetization and strain due to magnetic field and stress in magnetostrictive materials. The enthalpy of a ferromagnetic body with cubic symmetry is written with magnetization and strain as the internal states and the equilibrium strains are calculated by minimizing the enthalpy. Evaluating the enthalpy using the equilibrium strains, functions of the magnetization orientation, results in an enthalpy expression devoid of strain. By inspecting this expression, the magnetoelastic, elastic, and mechanical work energies are identified to be equivalent to the stress-induced anisotropy plus magnetostriction-induced fourth order anisotropy. It is shown that as long as the value of fourth order crystalline anisotropy constant $K_1$ includes the value of magnetostriction-induced fourth order anisotropy constant $\Delta K_1$, energy formulations involving magnetoelastic, elastic, and mechanical work energies are equivalent to those involving stress-induced anisotropy energy. Further, since the stress-induced anisotropy is only given for a uniaxial applied stress, an expression is developed for a general 3D stress.

Keywords: magnetostriction, magnetoelastic, magnetocrystalline, anisotropy, energy, delta K, 3-dimensional

1. INTRODUCTION

Magnetostriuctive materials exhibit dimensional and magnetization changes in response to magnetic fields and stresses. The dimensional change due to the application of a magnetic field is exploited for actuation. The magnetization change due to stress is used for sensing. The development of new magnetostriective materials like Iron-Gallium alloys (Galfenol) demonstrating moderate magnetostriction and steel-like structural properties enables these materials to be used in applications involving bending, torsion, etc. in 3D structures. Such applications require modeling tools that capture the nonlinear constitutive response of magnetostriective materials.

Nonlinear modeling of magnetostriiction often involves the calculation of the system’s free energy,\textsuperscript{1} where the internal states are magnetization and strain and the applied work is due to mechanical stresses and magnetic fields. Alternative energy approaches\textsuperscript{2–4} use only the magnetization as the internal state and incorporate magneto-mechanical coupling by including the anisotropy energy induced by applied stress in the system’s free energy. The energy with only the magnetization as an internal state is

$$E_T = E_K + E_{\sigma}^{\text{aniso}} - W_{\text{mag}},$$

where the first, second, and third terms are magnetocrystalline anisotropy, stress-induced anisotropy, and magnetic work (Zeeman energy) energies respectively. Although the second term in the energy expression is sometimes

\textsuperscript{1} Further author information: (Send correspondence to C.M)
C.M.: E-mail: chaitanya.mudivarthi@gmail.com, Phone: 1 301 405 2002, fax: 1 301 314 9001
A.B.F.: E-mail: aflatau@umd.edu, Phone: 1 301 405 1131
M.J.D.: E-mail: dapino.1@osu.edu, Phone: 1 614 688 3689
referred to as magnetoelastic energy, the works of Kittel et al. and Chikazumi et al. show that it is actually
the stress-induced anisotropy energy. The expression for the magnetoelastic energy involves strains as will be
described in equation (6) while the expression for stress-induced anisotropy does not.

Jiles and Thoelke\(^2\) used the energy expression (1) for modeling the effects of stress and anisotropy on the
magnetization and magnetostriction of Terfenol-D. The approach calculates the bulk magnetization and magne-
tostriction as the sum of the contributions from the various equilibria without explicitly calculating the fractional
occupancy of the material in each equilibrium. Armstrong\(^3\) used an empirical expression to explicitly calculate
the energy distribution and subsequently the bulk material magnetization and magnetostriction of Terfenol-D.
Evans and Dapino\(^7\) employed a thermodynamic framework to calculate the energy distribution and model the
magnetization and magnetostriction of Galfenol. The Armstrong model has also been used to model actuation
and sensing behavior of Iron-Gallium alloys.\(^8\)–\(^10\)

The modeling approaches discussed here agree well with experiments despite their use of an energy potential
lacking strain as an internal state. In this paper, the conditions are given for which (1) is equivalent to an energy
potential which includes strain through the elastic, magnetoelastic, and mechanical work energies. Under these
conditions, the accuracy of models utilizing (1) is not effected by the absence of strain. Further, the energy
expression (1) is typically for 3-D magnetic fields and unidirectional stresses. Therefore, an expression suitable
not only for 3-D magnetic fields but also for 3-D stresses is developed which is of importance for modeling 3-D
Galfenol behavior.

2. ENERGY DERIVATION

Most widely used magnetostrictive materials like Nickel, Terfenol-D and Galfenol have a cubic symmetry and
hence all the energy expressions used or derived in this section are for materials with cubic symmetry. As
discussed in the introduction, an energy expression commonly used in magnetomechanical models is the sum of
the magnetocrystalline anisotropy energy

\[
E_K = K_1 (\alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2) + K_2 (\alpha_1^2 \alpha_2^2 \alpha_3^2),
\]

stress-induced anisotropy energy

\[
E_{\text{anis}} = -\frac{3}{2} \lambda_{100} \sigma \left\{ \alpha_1^2 \beta_1^2 + \alpha_2^2 \beta_2^2 + \alpha_3^2 \beta_3^2 - \frac{1}{3} \right\} - 3 \lambda_{111} \sigma \left( \alpha_1 \alpha_2 \beta_1 \beta_2 + \alpha_2 \alpha_3 \beta_2 \beta_3 + \alpha_3 \alpha_1 \beta_3 \beta_1 \right),
\]

and magnetic work or Zeeman energy

\[
W_{\text{mag}} = \mu_0 M^T H.
\]

In the above equations, \(K_1\) and \(K_2\) are fourth and sixth order cubic anisotropy constants respectively and \(\lambda_{100}\)
and \(\lambda_{111}\) are the magnetostriction constants. The direction cosines \(\alpha_i\) of the magnetization vector \(M\) depend on
the applied magnetic field \(H\) and stress \(\sigma\) acting along a direction given by the direction cosines \(\beta_i\). Equilibrium
directions of the magnetization can be calculated by minimizing the total energy. However, this total energy, as
discussed in the introduction, lacks magnetoelastic \(E_{\text{magel}}\), elastic \(E_{\text{el}}\), and mechanical work \(W_{\text{mech}}\) terms.

To investigate the thermodynamic accuracy of using \(E_{\text{anis}}\) in place of \(E_{\text{magel}}, E_{\text{el}}, \) and \(W_{\text{mech}}\), the internal
energy and work energy of a ferromagnetic system are expressed as the sum of the \(E_K, E_{\text{magel}}, \) and \(E_{\text{el}}\) and sum
of the \(W_{\text{mag}}\) and \(W_{\text{mech}}\) respectively. The Gibbs free energy of the system is then formulated as the Legendre
transformation of the internal energy. Assuming isothermal and isentropic processes, the Gibbs free energy is
reduced to the enthalpy of the system

\[
\mathcal{H} = E_K + E_{\text{magel}} + E_{\text{el}} - W_{\text{mag}} - W_{\text{mech}}.
\]

The expression for \(E_K\) is given by equation (2) and the expression for the magnetoelastic energy, derived by Néel\(^5\)–\(^11\)
considering bond straining due to atomic magnetic moment rotation, is given by

\[
E_{\text{magel}} = B_1 \left\{ \epsilon_{\text{xx}} \left( \alpha_1^2 - \frac{1}{3} \right) + \epsilon_{\text{yy}} \left( \alpha_2^2 - \frac{1}{3} \right) + \epsilon_{\text{zz}} \left( \alpha_3^2 - \frac{1}{3} \right) \right\} + B_2 \left( \epsilon_{\text{xy}} \alpha_1 \alpha_2 + \epsilon_{\text{yz}} \alpha_2 \alpha_3 + \epsilon_{\text{zx}} \alpha_3 \alpha_1 \right)
\]

\[
= b \epsilon,
\]

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where
\[ b = \begin{bmatrix} B_1 (\alpha_1^2 - \frac{1}{3}) & B_1 (\alpha_2^2 - \frac{1}{3}) & B_1 (\alpha_3^2 - \frac{1}{3}) & B_2 \alpha_1 \alpha_2 & B_2 \alpha_2 \alpha_3 & B_2 \alpha_3 \alpha_1 \end{bmatrix} \] (8)
and
\[ \epsilon = \begin{bmatrix} \epsilon_{xx} & \epsilon_{yy} & \epsilon_{zz} & \epsilon_{xy} & \epsilon_{yz} & \epsilon_{zx} \end{bmatrix}^T. \] (9)

Constants \( B_1 \) and \( B_2 \) are called magnetomechanical coupling constants\(^6,12\) and can be calculated from the values of magnetostriction and elastic constants of a ferromagnetic material as will be seen later. The axis directions \( x, y, \) and \( z \) correspond to the \( <100> \) crystallographic directions of the material.

The elastic energy can be written as
\[ E_{el} = \frac{1}{2} \sigma^T \tilde{C} \epsilon, \] (10)
where \( \tilde{C} \) is the stiffness matrix. For cubic materials, \( \tilde{C} \) is expressed using elastic constants \( c_{11}, c_{12}, \) and \( c_{44} \) as
\[ \tilde{C} = \begin{bmatrix} c_{11} & c_{12} & c_{12} & 0 & 0 & 0 \\ c_{12} & c_{11} & c_{12} & 0 & 0 & 0 \\ c_{12} & c_{12} & c_{11} & 0 & 0 & 0 \\ 0 & 0 & 0 & c_{44} & 0 & 0 \\ 0 & 0 & 0 & 0 & c_{44} & 0 \\ 0 & 0 & 0 & 0 & 0 & c_{44} \end{bmatrix}. \] (11)

The expression for \( W_{mag} \) is given by equation (4), while the expression for \( W_{mech} \) can be written as the product of an externally applied stress \( (\sigma) \) given in equation (12) and the resulting strain \( (\epsilon) \).
\[ \sigma^T = \begin{bmatrix} \sigma_{xx} & \sigma_{yy} & \sigma_{zz} & \sigma_{xy} & \sigma_{yz} & \sigma_{zx} \end{bmatrix}. \] (12)
\[ W_{mech} = \sigma^T \epsilon \] (13)

Magnetization can be expressed as \( \mathbf{M} = M_s [\alpha_1 \alpha_2 \alpha_3]^T \), where \( M_s \) is the saturation magnetization, which is a material property. Using this, the enthalpy of the system can be written as a function of \( \alpha_i \) and \( \epsilon \). Therefore, the variables \( \alpha_i \) and \( \epsilon \) can now be termed as the system’s internal variables. Using equations (2), (4), (7), (10), and (13) the enthalpy can be expressed as
\[ \mathcal{H}(\alpha_i, \epsilon) = K_1 (\alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2) + K_2 (\alpha_2^2 \alpha_3^2 \alpha_1^2) + b \epsilon + \frac{1}{2} \epsilon^T \tilde{C} \epsilon - \mu_0 \mathbf{M}^T \mathbf{H} - \sigma^T \epsilon. \] (14)

The equilibrium states of the system can be calculated by minimizing \( \mathcal{H} \) with respect to its internal variables \( \alpha_i \) and \( \epsilon \). It is assumed that \( \mathcal{H}(\alpha_i, \epsilon) \) is a continuous function of \( \alpha_i \) and \( \epsilon \) and has continuous second order partial derivatives.

2.1 Equilibrium strains

Some of the earlier works\(^5,6,12,13\) that have derived the equilibrium strains did so under an assumption of a zero applied stress. In this work, the expression for the equilibrium strains is derived assuming a constant 3-D stress. The enthalpy of the system, which is defined in equation (14) with \( \alpha_i \) and \( \epsilon \) as the internal variables, is minimized with respect to \( \epsilon \)
\[ \frac{\partial \mathcal{H}(\alpha_i, \epsilon)}{\partial \epsilon} = 0, \] (15)
which gives
\[ \mathbf{b}^T + \tilde{C} \epsilon - \sigma = 0. \] (16)
Solving for \( \epsilon \) yields
\[ \epsilon^* = \tilde{C}^{-1} \sigma - \tilde{C}^{-1} \mathbf{b}^T = \epsilon_{mech} + \lambda. \] (17)
Taking a second partial derivative with respect to $\epsilon$ yields

$$\frac{\partial^2 H (\alpha_i, \epsilon)}{\partial \epsilon^2} = \tilde{C},$$

which is a positive value and hence $\epsilon^*$ corresponds to a relative minimum of $H$. Therefore, $\epsilon^*$ is the equilibrium strain.

It is noted that the equilibrium strain derived earlier for zero stress included only the second part of the equation (17), which is the magnetostrictive $\lambda = -\tilde{C}^{-1}b^T$ strain. For the non-zero stress condition derived here, the equilibrium strain is a superposition of purely mechanical $\epsilon_{\text{mech}} = C^{-1}\sigma$ and magnetostrictive strains.

Using equations (8) and (11), the magnetostrictive strain can be calculated as

$$\lambda = \left[ \frac{B_1}{c_{12} - c_{11}} (\alpha_1^2 - \frac{1}{3}) - \frac{B_1}{c_{12} - c_{11}} (\alpha_2^2 - \frac{1}{3}) - \frac{B_1}{c_{12} - c_{11}} (\alpha_3^2 - \frac{1}{3}) - \frac{B_2}{c_{44}} \alpha_1 \alpha_2 - \frac{B_2}{c_{44}} \alpha_2 \alpha_3 - \frac{B_2}{c_{44}} \alpha_3 \alpha_1 \right].$$

The elongation due to magnetostriction along any direction $(\gamma_1, \gamma_2, \gamma_3)$ can be evaluated using the expression

$$\frac{\delta l}{l} = \lambda_{xx} \gamma_1^2 + \lambda_{yy} \gamma_2^2 + \lambda_{zz} \gamma_3^2 + \lambda_{xy} \gamma_1 \gamma_2 + \lambda_{yz} \gamma_2 \gamma_3 + \lambda_{zx} \gamma_3 \gamma_1,$$

which becomes

$$\frac{\delta l}{l} = \frac{B_1}{c_{12} - c_{11}} \left( \gamma_1^2 \alpha_1^2 + \gamma_2^2 \alpha_2^2 + \gamma_3^2 \alpha_3^2 - \frac{1}{3} \right) - \frac{B_2}{c_{44}} (\gamma_1 \gamma_2 \alpha_1 \alpha_2 + \gamma_2 \gamma_3 \alpha_2 \alpha_3 + \gamma_3 \gamma_1 \alpha_3 \alpha_1)$$

by substituting the expressions for $\lambda_{ii}$ from equation (19).

The magnetostriction along the direction [100] occurs when the magnetization is along this direction. This can be calculated to be

$$\lambda_{100} = -\frac{2}{3} \frac{B_1}{c_{11} - c_{12}}$$

by using $\alpha_1 = \gamma_1 = 1$ and $\alpha_2 = \alpha_3 = \gamma_2 = \gamma_3 = 0$ in equation (21). Similarly, magnetostriction along [111] can be obtained to be

$$\lambda_{111} = -\frac{1}{3} \frac{B_2}{c_{44}}$$

by using $\alpha_i = \gamma_i = \frac{1}{\sqrt{3}}$ in equation (21).

Using equations (22) and (23), the magnetoelastic coupling constants can be expressed in terms of the magnetostrictive constants and elastic constants as

$$B_1 = -\frac{3}{2} \lambda_{100} (c_{11} - c_{12}); \quad B_2 = -3 \lambda_{111} c_{44}$$

(24)

### 2.2 Expression for enthalpy

The equilibrium strain $\epsilon^*$ in equation (17) can be substituted back into equation (14) to obtain an expression for the enthalpy in terms of $\alpha_i$. Minimizing this expression with respect to $\alpha_i$ will yield the equilibrium magnetization directions. As a result, any term that is not a function of $\alpha_i$ can be ignored.

$$H (\alpha_i) = K_1 (\alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2) + K_2 (\alpha_1^2 \alpha_2^2 \alpha_3^2) + E_{\text{magel}} (\epsilon^*) + E_{\text{cl}} (\epsilon^*) - W_{\text{mag}} - W_{\text{mech}} (\epsilon^*)$$

The expressions for $E_{\text{magel}} (\epsilon^*)$ and $E_{\text{cl}} (\epsilon^*) - W_{\text{mech}} (\epsilon^*)$ will be individually evaluated and then summed up together to get an expression for the enthalpy.
2.2.1 Expression for magnetoelastic energy

Substituting the equilibrium strain (\(\epsilon^*\)) in equation (6) and expanding gives the magnetoelastic energy as the sum of two terms:

\[
E_{magel}(\epsilon^*) = b(\epsilon_{mech} + \lambda) = b\epsilon_{mech} + b\lambda.
\]  

(26)

Supposing a stress (\(\sigma\)) acting on the ferromagnetic body along a direction (\(\beta_1, \beta_2, \beta_3\)), the corresponding mechanical strain \(\epsilon_{mech} = \epsilon_{ii}\) can be written as:

\[
\epsilon_{mech} = \sigma \left[ \frac{\beta_i^2}{c_{ii1} - c_{ii2}} + s_{12} \frac{\beta_j^2}{c_{ii1} - c_{ii2}} + s_{12} \frac{\beta_k^2}{c_{ii1} - c_{ii2}} + s_{12} \frac{\beta_j \beta_k}{c_{44}} \frac{\beta_j \beta_k}{c_{44}} \right]^T,
\]

(27)

where \(s_{12}\) is a compliance constant.

Substituting equation (27) in the first term of equation (26) and using equation (7) yields

\[
b\epsilon_{mech} = B_1 \left\{ e_{xx} \left( \frac{\alpha_i^2 - 1}{3} \right) + e_{yy} \left( \frac{\alpha_j^2 - 1}{3} \right) + e_{zz} \left( \frac{\alpha_k^2 - 1}{3} \right) \right\} + B_2 \left( e_{xy} \alpha_1 \alpha_2 + e_{yz} \alpha_2 \alpha_3 + e_{zx} \alpha_3 \alpha_1 \right)
\]

\[
= \frac{B_1}{c_{11} - c_{12}} \sigma \left\{ \frac{\alpha_i^2 \beta_1}{c_{11} - c_{12}} + \frac{\alpha_j^2 \beta_2}{c_{11} - c_{12}} + \frac{\alpha_k^2 \beta_3}{c_{11} - c_{12}} - \frac{1}{3} \right\} + \frac{B_2}{c_{44}} \sigma \left( \alpha_1 \alpha_2 \beta_1 \beta_2 + \alpha_2 \alpha_3 \beta_2 \beta_3 + \alpha_3 \alpha_1 \beta_3 \beta_1 \right),
\]

(28)

which can be simplified using the expressions for \(B_1\) and \(B_2\) from equation (24) to obtain

\[
b\epsilon_{mech} = -\frac{3}{2} \lambda_{100} \sigma \left\{ \frac{\alpha_i^2 \beta_1}{c_{11} - c_{12}} + \frac{\alpha_j^2 \beta_2}{c_{11} - c_{12}} + \frac{\alpha_k^2 \beta_3}{c_{11} - c_{12}} - \frac{1}{3} \right\} - 3\lambda_{111} \sigma \left( \alpha_1 \alpha_2 \beta_1 \beta_2 + \alpha_2 \alpha_3 \beta_2 \beta_3 + \alpha_3 \alpha_1 \beta_3 \beta_1 \right).
\]

(29)

The above expression is the same as the expression for the stress-induced anisotropy energy given in equation (3). Hence, the first term of equation (26) can be replaced with

\[
b\epsilon_{mech} = E_{aniso}^\sigma.
\]

(30)

The second term in equation (26) can be evaluated by substituting from equations (8) and (19) to obtain

\[
b\lambda = B_1 \left\{ \frac{\lambda_{xx}}{c_{11} - c_{12}} \left( \frac{\alpha_i^2 - 1}{3} \right) + \frac{\lambda_{yy}}{c_{11} - c_{12}} \left( \frac{\alpha_j^2 - 1}{3} \right) + \frac{\lambda_{zz}}{c_{11} - c_{12}} \left( \frac{\alpha_k^2 - 1}{3} \right) \right\} + B_2 \left( \lambda_{xy} \alpha_1 \alpha_2 + \lambda_{yz} \alpha_2 \alpha_3 + \lambda_{zx} \alpha_3 \alpha_1 \right)
\]

\[
= -\frac{B_1^2}{c_{11} - c_{12}} \left\{ \left( \frac{\alpha_i^2}{3} \right)^2 + \left( \frac{\alpha_j^2}{3} - 1 \right)^2 + \left( \frac{\alpha_k^2}{3} - 1 \right)^2 \right\} - \frac{B_2^2}{c_{44}} \left( \alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2 \right),
\]

(31)

which can be simplified by utilizing the expressions for \(B_1\) and \(B_2\) from equation (24) to

\[
b\lambda = \frac{9}{2} \left[ \lambda_{100}^2 (c_{11} - c_{12}) - 2\lambda_{111}^2 c_{44} \right] \left( \alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2 \right).
\]

(32)

Adding equations (30) and (32) gives the expression for the magnetoelastic energy

\[
E_{magel}(\epsilon^*) = \frac{9}{2} \left[ \lambda_{100}^2 (c_{11} - c_{12}) - 2\lambda_{111}^2 c_{44} \right] \left( \alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2 \right) + E_{aniso}^\sigma.
\]

(33)

It is noted that the magnetoelastic energy equals the stress-induced anisotropy energy plus an additional fourth order anisotropy energy term.
2.2.2 Expressions for elastic and mechanical work energies

The expression for $E_{el} (\epsilon^*) - W_{mech} (\epsilon^*)$ is evaluated by substituting the equilibrium strain ($\epsilon^*$) from equation (17) into equations (10) and (13) to obtain

$$E_{el} (\epsilon^*) - W_{mech} (\epsilon^*) = \frac{1}{2} (\epsilon_{mech} + \lambda)^T \tilde{C} (\epsilon_{mech} + \lambda),$$

which can be expanded and simplified to

$$E_{el} (\epsilon^*) - W_{mech} (\epsilon^*) = \frac{1}{2} \lambda^T \tilde{C} \lambda - \frac{1}{2} \epsilon_{mech}^T \tilde{C} \epsilon_{mech}. \tag{35}$$

The second term in the equation (35) is devoid of $\alpha_i$. Therefore, as mentioned before, the equilibrium $\alpha_i$ do not depend on this term and can be ignored.

The first term in the equation (35) can be expanded by using equation (11) to obtain

$$\frac{1}{2} \lambda^T \tilde{C} \lambda = \frac{1}{2} \left[ c_{11} (\lambda_{xx}^2 + \lambda_{yy}^2 + \lambda_{zz}^2) + c_{44} (\lambda_{xy}^2 + \lambda_{yz}^2 + \lambda_{zx}^2) + 2c_{12} (\lambda_{xz} \lambda_{yy} + \lambda_{yy} \lambda_{zz} + \lambda_{zz} \lambda_{xx}) \right]. \tag{36}$$

Substituting the values of $\lambda_{ii}$ from equation (19) gives

$$\frac{1}{2} \lambda^T \tilde{C} \lambda = \frac{1}{2} c_{11} \left( \sum_{i=1}^2 \frac{B_i^2}{c_{11} - c_{12}} \left( \left( \frac{\alpha_i^2 - 1}{3} \right)^2 + \left( \frac{\alpha_i^2 - 1}{3} \right)^2 + \left( \frac{\alpha_i^2 - 1}{3} \right)^2 \right) \right) + \cdots$$

$$\frac{c_{12}}{2 c_{44}} \left( \sum_{i=1}^2 \frac{B_i^2}{c_{11} - c_{12}} \left( \left( \frac{\alpha_i^2 - 1}{3} \right) \left( \frac{\alpha_i^2 - 1}{3} \right) + \left( \frac{\alpha_i^2 - 1}{3} \right) \left( \frac{\alpha_i^2 - 1}{3} \right) + \left( \frac{\alpha_i^2 - 1}{3} \right) \left( \frac{\alpha_i^2 - 1}{3} \right) \right) \right) + \cdots$$

$$\frac{1}{2} c_{44} \left( \sum_{i=1}^2 \frac{B_i^2}{c_{11} - c_{12}} \left( \alpha_i^2 \alpha_2 + \alpha_2^2 \alpha_3 + \alpha_3^2 \alpha_1 \right) \right), \tag{37}$$

which can be simplified by utilizing the expressions for $B_1$ and $B_2$ from equation (24) to yield

$$\frac{1}{2} \lambda^T \tilde{C} \lambda = -\frac{9}{4} \left[ \lambda_{100}^2 (c_{11} - c_{12}) - 2 \lambda_{111}^2 c_{44} \right] \left( \alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2 \right). \tag{38}$$

Therefore,

$$E_{el} (\epsilon^*) - W_{mech} (\epsilon^*) = -\frac{9}{4} \left[ \lambda_{100}^2 (c_{11} - c_{12}) - 2 \lambda_{111}^2 c_{44} \right] \left( \alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2 \right) \tag{39}$$

ignoring the energy term that is independent of $\alpha_i$.

Adding equations (33) and (39) gives

$$E_{magel} (\epsilon^*) + E_{el} (\epsilon^*) - W_{mech} (\epsilon^*) = \frac{9}{4} \left[ \lambda_{100}^2 (c_{11} - c_{12}) - 2 \lambda_{111}^2 c_{44} \right] \left( \alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2 \right) + E_{aniso}. \tag{40}$$

The first term in equation (40) is a contribution to the anisotropy energy due to the equilibrium magnetostrictive strains. A fourth-order magnetostrictive anisotropy constant

$$\Delta K_1 = \frac{9}{4} \left[ \lambda_{100}^2 (c_{11} - c_{12}) - 2 \lambda_{111}^2 c_{44} \right] \tag{41}$$

is defined and equation (40) becomes

$$E_{magel} (\epsilon^*) + E_{el} (\epsilon^*) - W_{mech} (\epsilon^*) = \Delta K_1 \left( \alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2 \right) + E_{aniso}. \tag{42}$$

Substituting equation (42) in equation (25) yields the expression for the enthalpy of the system

$$H (\alpha_i) = (K_1 + \Delta K_1) \left( \alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2 \right) + K_2 \left( \alpha_1^2 \alpha_2^2 \alpha_3^2 \right) + E_{aniso} - W_{mag}. \tag{43}$$
2.3 Discussion
Comparing equation (43) to the total energy expression in equation (1), it can be seen that the only difference between the two is the fourth order anisotropy constant $K_1$. The total energy in equation (1) uses the magnetocrystalline anisotropy energy with fourth order anisotropy constant as $K_1$, whereas the enthalpy expression derived here shows that it is in fact $K_1$ plus the magnetostriction-induced fourth order anisotropy constant $\Delta K_1$. Kittel et al.\textsuperscript{6} described that an experimental determination of $K_1$ can be made only if the ferromagnetic body is held at a constant strain. If the ferromagnetic body is allowed to strain, then the value observed for $K_1$ would in fact be $K_1 + \Delta K_1$. Since it is experimentally difficult to maintain a constant strain, the value of theoretical $K_1$ is usually obtained by subtracting the calculated value of $\Delta K_1$ from the experimentally observed $K_1$ value.

The values of the fourth order cubic anisotropy constant $K_1$ for FeGa alloys reported by Rafique et al.\textsuperscript{14} do not include the corrective $\Delta K_1$ term. Therefore, the reported values are actually the values of $K_1 + \Delta K_1$. The appropriate values for $K_1$ can be calculated by subtracting $\Delta K_1$ calculated using equation (41) with values of the magnetostriction and elastic constants.

Figure 1 shows the corrected or theoretical $K_1$ values for different compositions of FeGa along with the reported values from Rafique et al.\textsuperscript{14} and calculated values of $\Delta K_1$. For the calculation of $\Delta K_1$, magnetostrictive values reported by Clark et al.\textsuperscript{15} and stiffness constant values reported by Wuttig et al.\textsuperscript{16} were used.

![Figure 1. Experimentally determined (uncorrected) $K_1$ and Corrected $K_1$ for different at% Ga FeGa alloys](image1.png)

![Figure 2. Percent deviation of uncorrected ($K_1 + \Delta K_1$) values from corrected values ($K_1$)](image2.png)

In figures 1 and 2 the deviation of corrected from uncorrected values of $K_1$ for FeGa alloys with less than 18\% Ga content, is less than 10\%, while close to 20\% Ga content, where $K_1$ becomes very small, it can be as high as 200\%. However, since $K_1$ for FeGa alloys around 20 at\% Ga is very small, crystalline anisotropy energy is dominated by the other energy terms and as a result, the usage of $K_1$ instead of $K_1 + \Delta K_1$ will only affect very low magnetic field or mechanical stress conditions. Moreover, the percentage deviations are within the error bounds of measured $K_1$ values (uncorrected) reported by Rafique et al. Therefore, it is concluded that no changes are required for the reported values of $K_1$ for FeGa alloys. In magnetostrictive materials other than FeGa, $\Delta K_1$ may be more significant. For thermodynamic consistency, when $E_{\text{magg}}, E_{\text{el}},$ and $W_{\text{mech}}$ is replaced with $E_{\sigma}^{\text{miso}}$ in the total energy, the total anisotropy constant $K_1 + \Delta K_1$ should be used.

2.4 Energy expression for 3D stresses
For 3-D modeling, the energy expression in equation (43) needs slight modification. While equation (43) describes the free energy expression for 3-D magnetic fields, it is valid only for unidirectional stresses. Equation (43) can be easily extended to incorporate 3-D stresses.
It is well known that any 3-D stress tensor acting on a body can be decomposed into principal stresses and principle directions by solving the eigenvalue problem

\[
\begin{bmatrix}
\sigma_{xx} & \sigma_{xy} & \sigma_{xz} \\
\sigma_{xy} & \sigma_{yy} & \sigma_{yz} \\
\sigma_{xz} & \sigma_{yz} & \sigma_{zz}
\end{bmatrix}
\begin{bmatrix}
\beta_1 \\
\beta_2 \\
\beta_3
\end{bmatrix}
= \sigma
\begin{bmatrix}
\beta_1 \\
\beta_2 \\
\beta_3
\end{bmatrix},
\]

which results in three principle stresses (eigenvalues), \(\sigma_j\) acting along three principle directions (eigenvectors), \((\beta_{1j}, \beta_{2j}, \beta_{3j})\).

The mechanical strain resulting from all the three principle stresses can be evaluated by the principle of superposition

\[
\epsilon_{mech} = \sum_{j=1}^{3} \sigma \left[ \frac{\beta_{1j}^2}{c_{11}-c_{12}} + \frac{\beta_{2j}^2}{c_{11}-c_{12}} + \frac{\beta_{3j}^2}{c_{11}-c_{12}} + \frac{\beta_{1j}\beta_{2j}}{c_{44}} \right].
\]

From equations (30) and (45), it directly follows that the net stress-induced anisotropy energy is sum of the stress induced anisotropy energies due to each principle stress

\[
E_{aniso}^{3D} = \sum_{j=1}^{3} \frac{3}{2} \lambda_{100} \sigma \left\{ \alpha_1^2 \beta_{1j}^2 + \alpha_2^2 \beta_{2j}^2 + \alpha_3^2 \beta_{3j}^2 \right\} - \frac{1}{3} \lambda_{111} \sigma \left( \alpha_1 \alpha_2 \beta_{1j} \beta_{2j} + \alpha_2 \alpha_3 \beta_{2j} \beta_{3j} + \alpha_3 \alpha_1 \beta_{3j} \beta_{1j} \right)
\]

\[
= \sum_{j=1}^{3} E_{\sigma}^{aniso} (\sigma_j, \beta_{ij}).
\]

Therefore, the enthalpy expression that must be used to model the response of ferromagnetic materials subjected to 3-D magnetic fields and mechanical stresses is given by

\[
H(\alpha_i) = (K_1 + \Delta K_1) \left( \alpha_1^2 \alpha_2^2 + \alpha_2^2 \alpha_3^2 + \alpha_3^2 \alpha_1^2 \right) + K_2 \left( \alpha_1^2 \alpha_2^2 \alpha_3^2 \right) + \sum_{j=1}^{3} E_{\sigma}^{aniso} (\sigma_j, \beta_{ij}) - W_{mag}.
\]

### 3. CONCLUSIONS

An expression for the enthalpy of a ferromagnetic body has been derived. Starting from the enthalpy expression, which includes the energy density due to magnetocrystalline anisotropy, magnetoelastic coupling, and elastic strain and work energy densities due to magnetic fields and mechanical stresses, the equilibrium strains were derived. It has been shown that, as expected, the equilibrium strain is the superposition of pure mechanical strain and pure magnetostrictive strain. Evaluating the enthalpy expression with the equilibrium strains yields an expression which has only the magnetization orientation as the internal state. This expression is interpreted as the sum of the magnetocrystalline anisotropy, the stress-induced anisotropy, and the magnetic work energies where the fourth-order magnetocrystalline anisotropy constant is the sum of the intrinsic magnetic constant \(K_1\) and an anisotropy change \(\Delta K_1\) due to magnetoelastic coupling. Since the experimental determination of the crystalline anisotropy constant is usually done at constant stress, reported values include \(\Delta K_1\). It was shown that the difference in \(K_1\) and \(K_1 + \Delta K_1\) is within the experimental error of measurement for reported values of FeGa. It was concluded that modeling approaches that use stress-induced anisotropy energy in place of elastic, magnetoelastic, and mechanical work energies are thermodynamically consistent as long the magnetocrystalline anisotropy coefficients include \(\Delta K_1\) due to magnetoelastic coupling. Further, an expression for enthalpy suitable for modeling the response of ferromagnetic materials subjected to 3-D magnetic fields as well as 3-D mechanical stresses was provided.

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