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Supplemental Information

Soft Matter Mechanics and the Mechanisms

Underpinning the Infrared Vision of Snakes

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First and Second Laws of Thermodynamics

Defining the dipole density per unit volume in the reference configuration by nominal polarization \mathbf{P} , the free energy density by $\Psi(\mathbf{F}, \mathbf{P}, T)$, and the local electric field by

$$\mathcal{E}[\mathbf{x}, \mathbf{P}] = \int_{\Omega_t} \frac{\varepsilon_0}{2} |\nabla_{\mathbf{x}} \xi|^2 = \int_{\Omega_R} \frac{\varepsilon_0}{2} J |\mathbf{F}^{-T} \nabla \xi|^2, \quad (1)$$

the total free energy of the body is:

$$A[\mathbf{x}, \mathbf{P}, T] = \int_{\Omega_R} \Psi(\mathbf{F}, \mathbf{P}, T) + \mathcal{E}[\mathbf{x}, \mathbf{P}]. \quad (2)$$

And the total internal energy of the body becomes

$$U[\mathbf{x}, \mathbf{P}, T] = \int_{\Omega_R} TS(\mathbf{F}, \mathbf{P}, T) + A[\mathbf{x}, \mathbf{P}, T], \quad (3)$$

where S is the local entropy density per unit volume in the reference configuration. We remark that here we have assumed that the body is homogeneous in space and time so Ψ , Φ , and S do not explicitly depend on \mathbf{X} . Considering Dirichlet boundary conditions

$$\begin{cases} \mathbf{x}(\mathbf{X}, t) = \mathbf{X} + \mathbf{u}_b(\mathbf{X}, t) & \text{on } \partial\Omega_R, \\ \xi(\mathbf{X}, t) = \xi_b(\mathbf{X}, t) & \text{on } \partial\Omega_R, \\ T(\mathbf{X}, t) = T_b(\mathbf{X}, t) & \text{on } \partial\Omega_R, \end{cases} \quad (4)$$

where \mathbf{u}_b , ξ_b , and T_b are the prescribed boundary displacement, prescribed electric potential, and prescribed temperature accordingly. The rate of work done to the system with no body force is

$$P_{\text{ext}} = \int_{\partial\Omega_R} (\dot{\mathbf{u}}_b \cdot \tilde{\mathbf{t}}^e - \xi_b \mathbf{D} \cdot \mathbf{n}), \quad (5)$$

where $\tilde{\mathbf{t}}^e$ is the traction applied to the system and $\mathbf{D} = -\varepsilon_0 J \mathbf{C}^{-1} \nabla \xi + \mathbf{F}^{-1} \mathbf{P}$ is the nominal electric displacement. The rate of total heat transferred into the body is

$$\dot{Q} = \int_{\Omega_R} r_e - \int_{\partial\Omega_R} \mathbf{q} \cdot \mathbf{n}, \quad (6)$$

where \mathbf{q} is the thermal flux and r_e is the rate of heat supply. Assuming that the considered system is stationary and neglect kinetic energy, first and second laws of thermodynamic imply that:

$$\begin{aligned} \frac{d}{dt}U &= P_{\text{ext}} + \dot{Q}, \\ \frac{d}{dt} \int_P S &\geq - \int_{\partial P} \frac{\mathbf{q} \cdot \mathbf{n}}{T} + \int_P \frac{r_e}{T}, \end{aligned} \quad (7)$$

where $P \subset \Omega_R$ is any material volume element in the reference configuration. With some tedious calculations addressed in Darbaniyan *et al.*,¹ we have

$$\frac{d}{dt} \mathcal{E} = \int_{\Omega_R} (\nabla \dot{\mathbf{x}} : \boldsymbol{\Sigma}_{\text{MW}} + \nabla \xi \cdot \mathbf{F}^{-1} \dot{\mathbf{P}}) - \int_{\partial \Omega_R} \xi_b \dot{\mathbf{D}} \cdot \mathbf{n}. \quad (8)$$

Where $\boldsymbol{\Sigma}_{\text{MW}} = (\mathbf{F}^{-T} \nabla \xi) \otimes (\varepsilon_0 J \mathbf{C}^{-1} \nabla \xi - \mathbf{F}^{-1} \mathbf{P}) - \frac{\varepsilon_0}{2} J |\mathbf{F}^{-T} \nabla \xi|^2 \mathbf{F}^{-T}$ is the so called the Maxwell stress. Substituting in first and second laws of thermodynamic and rearranging, the standard Coleman-Noll approach implies that

$$\begin{cases} \frac{\partial \Psi}{\partial \mathbf{P}} = -\mathbf{F}^{-T} \nabla \xi & \text{in } \Omega_R, \\ S(\mathbf{F}, \mathbf{P}, T) = -\frac{\partial}{\partial T} \Psi(\mathbf{F}, \mathbf{P}, T) & \text{in } \Omega_R, \\ -T \frac{d}{dt} \frac{\partial \Psi}{\partial T} - r_e + \nabla \cdot \mathbf{q} = 0 & \text{in } \Omega_R, \\ \nabla \cdot \boldsymbol{\sigma} = 0 & \text{in } \Omega_R, \\ \boldsymbol{\sigma} \cdot \mathbf{n} = \tilde{\mathbf{t}}^e & \text{on } \partial \Omega_R. \end{cases} \quad (9)$$

Where $\boldsymbol{\sigma} = \frac{\partial \Psi}{\partial \mathbf{F}} + \boldsymbol{\Sigma}_{\text{MW}}$ is the total stress applied to the system and with the assumption of the linear Fourier's law: $\mathbf{q} = -k \nabla T$, with $k \geq 0$.

Constitutive Relations

In this section we specify free energy function $\Psi(\mathbf{F}, \mathbf{P}, T)$ that conserves the principle of frame indifference and material symmetry. If $SO(3)$ be the group including all rigid rotations and $\mathcal{G} \subset SO(3)$ be the subgroup associated with material symmetry, for any $\mathbf{R} \in SO(3)$ and $\mathbf{Q} \in \mathcal{G}$, we should have

$$\Psi(\mathbf{R}\mathbf{F}, \mathbf{R}\mathbf{P}, T) = \Psi(\mathbf{F}, \mathbf{P}, T), \quad \Psi(\mathbf{F}\mathbf{Q}, \mathbf{P}, T) = \Psi(\mathbf{F}, \mathbf{P}, T). \quad (10)$$

Choosing the reference configuration as the equilibrium one with $(\mathbf{P}, T) = (0, T_0)$ in the absence of external loadings, we have $\Psi(\mathbf{F}, \mathbf{P}, T_0) \geq \Psi(\mathbf{I}, \mathbf{0}, T_0)$, with \mathbf{I} being the identity matrix. We assume that the material is isotropic with permittivity ε independent of temperature and deformation and incompressible at constant temperature. With the latter

assumption, thermoelastic incompressibility, the volume change caused by thermal expansion has to be taken into account.² To that end having α as the linear thermal expansion coefficient and $\Delta T = T - T_0$, we assume that:

$$J = 1 + 3\alpha\Delta T. \quad (11)$$

Considering Ψ_0 as a constant identified as the free energy density at $(\mathbf{F}, \mathbf{P}, T) = (\mathbf{I}, 0, T_0)$, one possible form for free energy density can be

$$\Psi(\mathbf{F}, \mathbf{P}, T) = \Psi_0 - CT \log \frac{T}{T_0} + \frac{\mu}{2}(\mathbf{I}_1 - 3) - \Pi|J - (1 + 3\alpha\Delta T)| + \frac{|\mathbf{P}|^2}{2J(\varepsilon - \varepsilon_0)}, \quad (12)$$

where Π is the Lagrange multiplier and ε_0 is the vacuum electric permittivity. From Eq.(9), we get:

$$\begin{cases} \nabla \cdot [\mu\mathbf{F} - \Pi J\mathbf{F}^{-T} - \frac{\varepsilon}{2}J|\mathbf{F}^{-T}\nabla\xi|^2\mathbf{F}^{-T} + \varepsilon J(\mathbf{F}^{-T}\nabla\xi) \otimes (\mathbf{C}^{-1}\nabla\xi)] = 0 & \text{in } \Omega_R, \\ [\mu\mathbf{F} - \Pi J\mathbf{F}^{-T} - \frac{\varepsilon}{2}J|\mathbf{F}^{-T}\nabla\xi|^2\mathbf{F}^{-T} + \varepsilon J(\mathbf{F}^{-T}\nabla\xi) \otimes (\mathbf{C}^{-1}\nabla\xi)] \cdot \mathbf{n} = \tilde{\mathbf{t}}^e & \text{on } \partial\Omega_R, \\ CT\dot{+} + 3\alpha T\dot{\Pi} = r_e + \nabla \cdot (k\nabla T) & \text{in } \Omega_R. \end{cases} \quad (13)$$

References

¹ F. Darbaniyan, K. Dayal, L. Liu, P. Sharma, Designing soft piezoelectric and electrocaloric materials using electrets, *Soft matter*.

² K. Volokh, *Mechanics of soft materials*, Springer, 2016.