Thermodynamics, plasticity and rheology

Ván P.

RMKI, Dep. Theor. Phys., Budapest, Hungary Montavid Research Group, Budapest, Hungary

- Minimal, simple small strain viscoelastoplasticity

-Minimal – reduction of empirical freedom

plasticity, rheology

- Simple – differential equation (1D, homogeneous)

Classical plasticity

1. Elastic stress strain relations 2. Plastic strain: $\varepsilon = \varepsilon^e + \varepsilon^p$ 3. Yield condition -f: $f(\sigma,...) = 0$ $(\dot{f}(\sigma,...) = 0)$ 4. Plastic potential -g: $\dot{\varepsilon}^p = \lambda \frac{\partial g}{\partial \sigma}(\sigma,...)$ (non-associative) 5. Evolution of plastic strain (non-ideal)

 \rightarrow Three empirical functions:

static potential *s*, yield function *f*, plastic potential $g \rightarrow$ Second Law?



Thermodynamic plasticity (Ziegler, Maugin, ..., Bazant, Houlsby,...)

1. Thermostatics – elastic stress strain relations 2. Plastic strain is a special internal variable 3. Dissipation potential = plastic potential 4. First order homogeneity $F = v \frac{\partial \Phi}{\partial \dot{\epsilon}^{p}}$

- → Relation to Second Law is established, evolution of plastic strain is derived.
- \rightarrow Two empirical functions:

static potential s, dissipation potential Φ

Thermodynamic rheology (Verhás, 1997) $s(e, \mathbf{F}, \boldsymbol{\xi}) = s_0(e, \mathbf{F}) - \frac{1}{2}\boldsymbol{\xi} : \boldsymbol{\xi}$ entropy

Entropy balance:

$$\rho \dot{s} + \nabla \cdot \mathbf{j}_{s} = \sigma_{s} = \mathbf{j}_{q} \cdot \nabla \frac{1}{T} + \frac{1}{T} \left(\mathbf{T} + \rho T \mathbf{F} \frac{\partial s}{\partial \mathbf{F}} \right) : \left(\dot{\mathbf{F}} \mathbf{F}^{-1} \right) - \frac{\rho}{T} \boldsymbol{\xi} : \dot{\boldsymbol{\xi}} \ge 0,$$

$$TP_{s} = \left(\boldsymbol{\sigma} + \rho \frac{\partial s}{\partial \boldsymbol{\varepsilon}}\right) : \dot{\boldsymbol{\varepsilon}} - \rho \boldsymbol{\xi} : \dot{\boldsymbol{\xi}} \ge 0,$$

isothermal, small strain

	Force	Flux – material function
Mechanical	$(\nabla \circ \mathbf{v}) = \dot{\mathbf{\varepsilon}}$	$\mathbf{\sigma} + \rho \frac{\partial s}{\partial \mathbf{\epsilon}}$
Rheological	$- ho\xi$	$\dot{\boldsymbol{\xi}} = \mathbf{G}(\boldsymbol{\varepsilon}, \dot{\boldsymbol{\varepsilon}}, \boldsymbol{\xi})$

Min?

Linear relations:

$$\begin{aligned} \mathbf{\sigma}^{v} = \mathbf{\sigma} + \rho \frac{\partial s}{\partial \mathbf{\epsilon}} \\ \dot{\mathbf{\xi}} \end{aligned} = \mathbf{L} \begin{pmatrix} \dot{\mathbf{\epsilon}} \\ -\rho \mathbf{\xi} \end{pmatrix} = \dots = \begin{pmatrix} l_{1} & l_{12} \\ l_{21} & l_{2} \end{pmatrix} \begin{pmatrix} \dot{\mathbf{\epsilon}} \\ -\rho \mathbf{\xi} \end{pmatrix}, \\ \text{isotropy, one dimension, } \dots \end{aligned}$$

Ideal elastic: $\rho \frac{\partial s}{\partial \varepsilon} = \rho \frac{\partial}{\partial \varepsilon} (-\mu \varepsilon : \varepsilon) = -2G\varepsilon,$
Second Law: $l_{1} > 0, \ l_{2} > 0, \ l_{1}l_{2} - l_{12}l_{21} > 0.$
The internal variable can (must?) be eliminated:
$$\mathbf{\sigma} + \tau \, \dot{\mathbf{\sigma}} = 2\eta \tau_{d} \ddot{\varepsilon} + 2\eta \dot{\varepsilon} + 2G\varepsilon \qquad \text{Jeffreys body}$$

relaxation material inertia creep
 $\tau = \frac{\rho}{l_{2}} > 0, \ \tau_{d} = \frac{\rho l_{1}}{2\eta l_{2}} > 0, \ 2\eta = \frac{l_{1}l_{2} - l_{12}l_{21} + 2G\rho}{l_{2}} > 0. \end{aligned}$

Further motivation:

- Rheology (creep and relaxation)+plasticity
- Non associated flow soils

dissipation potentials = normality = (nonlinear) Onsagerian symmetry ? Truesdell + dual variables (JNET, 2008, **33**, 235-254.) rheological thermodynamics (KgKK, 2008, **6**, 51-92) Is the thermodynamic framework too strong??

 \rightarrow computational and numerical stability

min+min=simple?

Unification:

dissipation potential – constitutive relations



simple

Strain hardening:

 $G=1, \ \overline{G}=0.05, \ l_1=0.001, \ l=0.00005; \ \sigma_{cr}=0.5, \ v=1$



Loading rate dependence:



v =10, 15, 25

simple

Creep:

 $G = 1, \ \overline{G} = 0.15, \ v_1 = 0.1, \ v_2 = 10.$



like Armstrong-Frederic kinematic hardening

simple

Hysteresis, Bauschinger effect, ratcheting:

 $G = 1, \ \overline{G} = 0.15, \ l_1 = 0.001, \ l = 0.001; \ \sigma_{cr} = 0.5, \ \sigma_{Max} = 0.6, \ v = 1$



Non-associative, Mohr-Coulomb, dilatancy

(according to Houlsby and Puzrin):

$$s(e, \mathbf{\epsilon}, \mathbf{\epsilon}^{p}) = s_{0}(e) - \frac{E}{2} (\varepsilon_{1} - b \varepsilon_{2}^{p})^{2} - \frac{G}{2} (\varepsilon_{2} - \varepsilon_{2}^{p})^{2} - \frac{\overline{G}}{2} \mathbf{\epsilon}^{p^{2}}$$

 $K = 1, G = 0.5, \overline{G} = 0, l_1 = 0.00005, l = 0.001; \sigma_{cr} = 0.5, b = -0.2, v = 1$



Discussion

- 1. Minimal viscoelastoplasticity
- 2. Possible generalizations
 - finite strain,
 - objective derivatives,
 - gradient effects,
 - modified Onsagerian coefficients,
 - yield beyond Tresca, etc...
- 3. Non-associative? + thermo
- 4. More general constitutive relations (gyroscopic forces?) + thermo
- 5. Differential equation (can be incremental)
- 6. Numerical stability (thermo)

Thank you for your attention!

Interlude – damping and friction

Dissipation in contact mechanics:

$$E(x, \dot{x}) = \frac{m\dot{x}^{2}}{2} + V(x)$$

$$0 \ge -D = \dot{E}(x, \dot{x}) = (m\ddot{x} - F)\dot{x} = S\dot{x}$$

$$S = S_{d} + S_{f} = -\beta\dot{x} - \mu N \frac{\dot{x}}{|\dot{x}|} \implies D = \beta \frac{\dot{x}^{2}}{2} + \mu N |\dot{x}|$$
Mechanics – constant force $F = \pm \mu N$
Plasticity:
$$S = -\beta (1 + \frac{|S|}{2}) \dot{x} \implies \dot{x} = -\beta^{-1}S$$

$$S = -\rho \left(\frac{1 + \mu N}{\mu N} \right) \xrightarrow{X} \Rightarrow X = \frac{1}{1 + \frac{|S|}{\mu N}} \longrightarrow -\mu N \le S \le \mu N$$
(spec. $S \parallel \dot{x}$)

Thermodinamics - Mechanics

