

Spring, 2022

ME 597 – Solid Mechanics II

Lecture 19 Review for the exam

News: Ready for the exam?



Mechanical Engineering

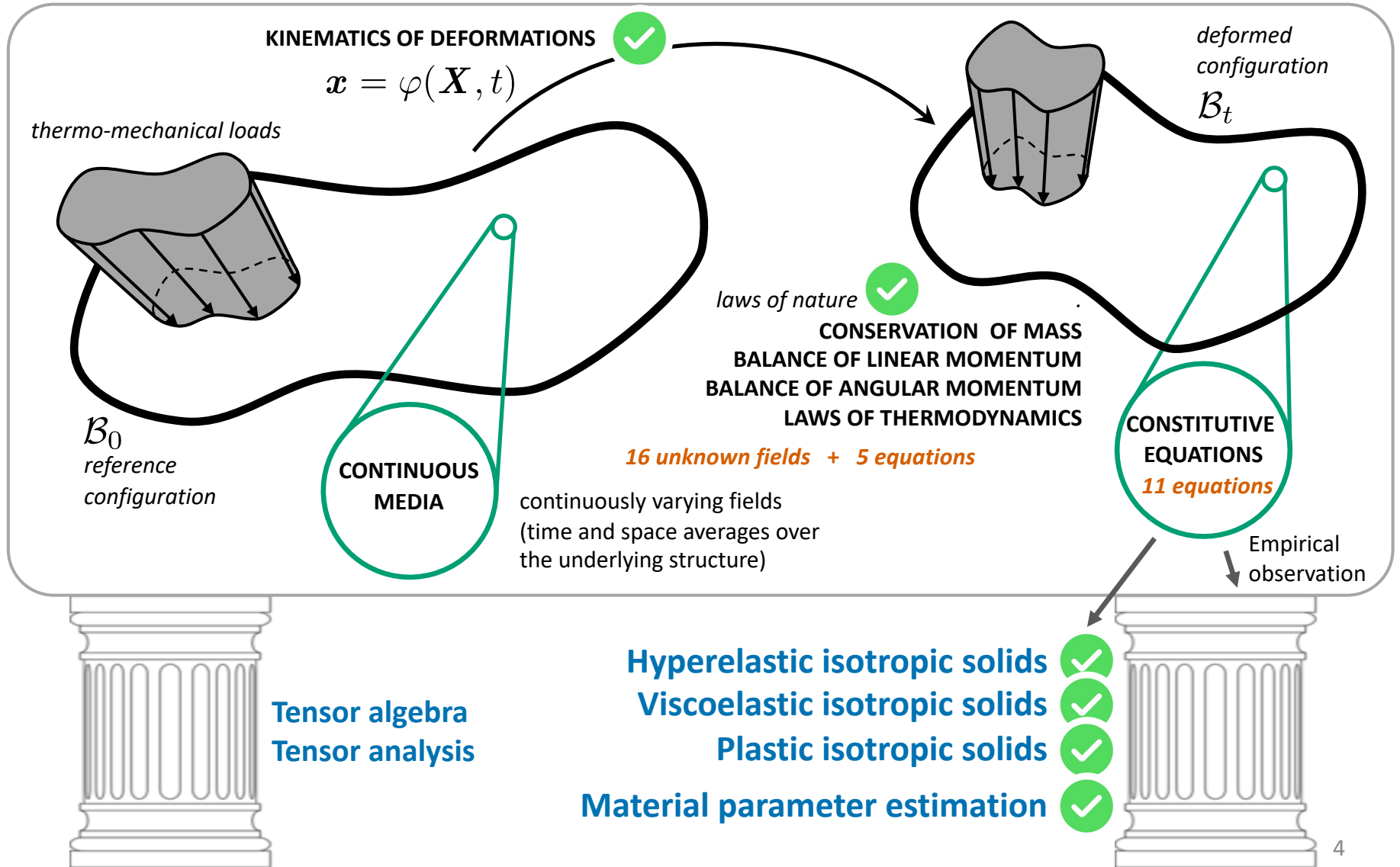
Instructor: Prof. Marcial Gonzalez

Last modified: 3/24/22 10:28:06 AM

ME 597 – Course outline - Syllabus

- Tensor algebra and tensor analysis (Lecture 2)
- Kinematics of deformations (Lecture 3-4)
- Mechanical conservation and balance laws (Lecture 5)
- Thermodynamics (Lecture 6)
- Constitutive relations (Lecture 7)
- Hyperelastic solids (Lecture 8-9)
- Viscoelastic solids (Lecture 10-11)
- Material parameter estimation (Lecture 12)
- Plasticity (Lecture 13-15)
- Homework set 1, 2, 3 & 4

ME 597 – Course outline - Syllabus



Introduction to tensors and vectors

Tensor algebra and analysis in Cartesian coordinates

$$B_{ij} = A_{ji} \iff \mathbf{B} = \mathbf{A}^T$$

$$D_{ijk} = A_{ij}v_k \iff \mathbf{D} = \mathbf{A} \otimes \mathbf{v} \qquad \mathbf{A} : \mathbf{B} = A_{ij}B_{ij}$$

$$v_i = A_{ik}u_k \iff \mathbf{v} = \mathbf{A}\mathbf{u}$$

1-order tensor (vector)

$$v_i \equiv \mathbf{v}[\mathbf{e}_i] \quad \mathbf{v} = v_i \mathbf{e}_i$$

2-order tensor

$$A_{ij} \equiv \mathbf{T}[\mathbf{e}_i, \mathbf{e}_j] \quad \mathbf{A} = A_{ij}(\mathbf{e}_i \otimes \mathbf{e}_j)$$

Symmetric, positive-definite 2-order tensor

$$\lambda_\alpha^{\mathbf{S}} \in \mathbb{R} \quad , \quad \boldsymbol{\Lambda}_\alpha^{\mathbf{S}} \cdot \boldsymbol{\Lambda}_\beta^{\mathbf{S}} = \delta_{\alpha\beta}$$

$$\mathbf{S} = \sum_{\alpha=1}^3 \lambda_\alpha^{\mathbf{S}} \boldsymbol{\Lambda}_\alpha^{\mathbf{S}} \otimes \boldsymbol{\Lambda}_\alpha^{\mathbf{S}} = S_{ij}(\mathbf{e}_i \otimes \mathbf{e}_j) \qquad \sqrt{\mathbf{S}} = \sum_{\alpha=1}^3 \sqrt{\lambda_\alpha^{\mathbf{S}}}(\boldsymbol{\Lambda}_\alpha^{\mathbf{S}} \otimes \boldsymbol{\Lambda}_\alpha^{\mathbf{S}})$$

- Gradient

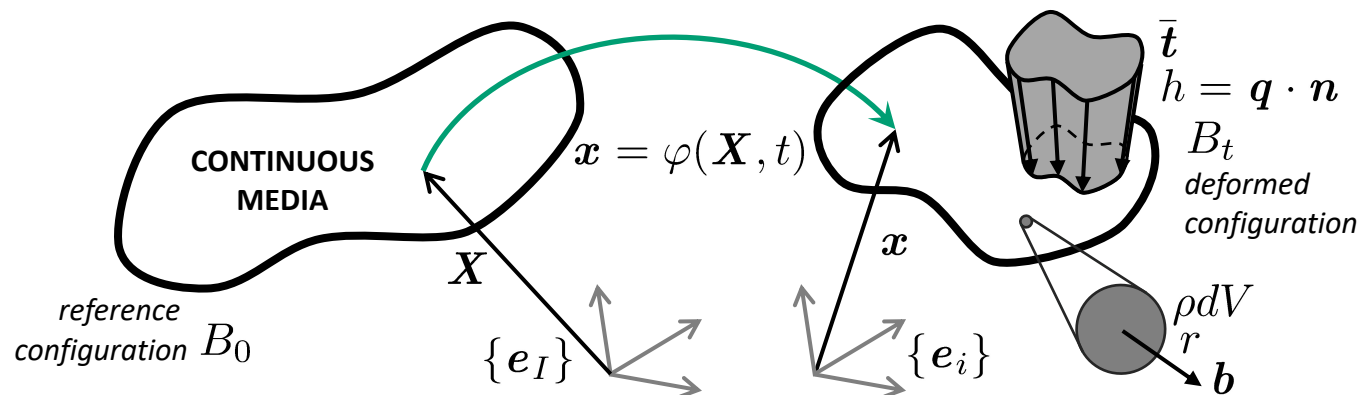
$$\nabla \mathbf{B} = \frac{\partial \mathbf{B}(\mathbf{x})}{\partial x_i} \otimes \mathbf{e}_i \qquad \nabla s = \frac{\partial s(\mathbf{x})}{\partial x_i} \mathbf{e}_i$$

- Divergence

$$\operatorname{div} \mathbf{B} = \frac{\partial \mathbf{B}(\mathbf{x})}{\partial x_i} \mathbf{e}_i \qquad \operatorname{div} \mathbf{v} = \frac{\partial v(\mathbf{x})}{\partial x_i} \cdot \mathbf{e}_i$$

Kinematics of deformations

- Deformation mapping $\mathbf{x} = \boldsymbol{\varphi}(\mathbf{X}, t)$
- Deformation gradient $\mathbf{F} = F_{iJ}(\mathbf{e}_i \otimes \mathbf{e}_J) = \frac{\partial x_i}{\partial X_J}(\mathbf{e}_i \otimes \mathbf{e}_J)$
- Right Cauchy-Green deformation tensor $\mathbf{C} = \mathbf{F}^T \mathbf{F}$
- Left Cauchy-Green deformation tensor $\mathbf{B} = \mathbf{F} \mathbf{F}^T$
- Jacobian $J = \det \mathbf{F}$ - Nanson's eqn. $\mathbf{n} dA = J \mathbf{F}^{-T} \mathbf{N} dA_0$
- Lagrangian strain tensor $\mathbf{E} = \frac{1}{2}(\mathbf{C} - \mathbf{I})$
- Material tensors, spatial tensors, two-point tensors.



Kinematics of deformations

Right and left polar decomposition $F = RU = VR$ $U = \sqrt{C}$

Q: How to calculate U or V ?

A: Using the spectral decomposition...

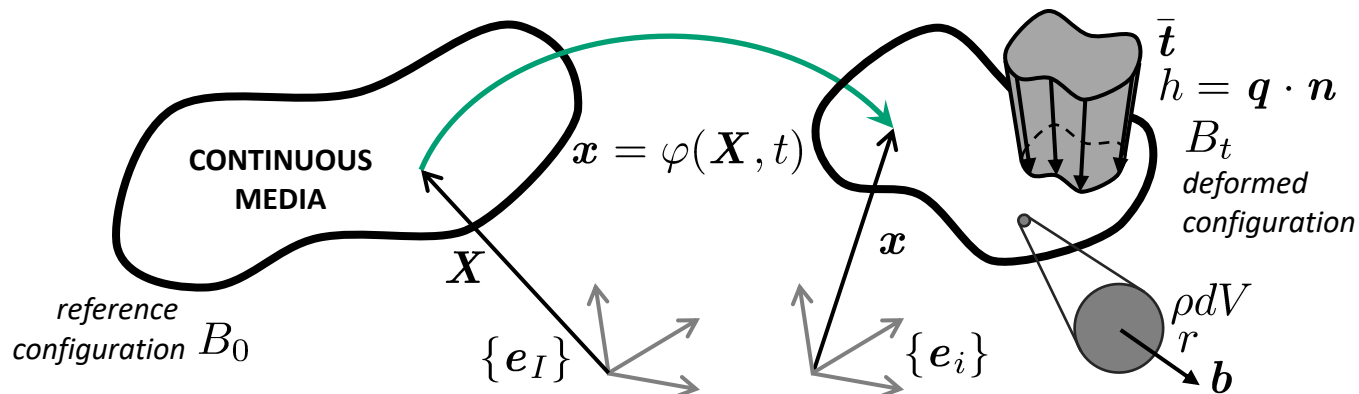
Q: How to calculate R ? A: Using $R = FU^{-1}$

Linear parts of kinematic fields

- Examples (for $X \rightarrow X + u(X)$, i.e., $F = I, \nabla_0 u = \nabla u$):
[the linearization is evaluated in the undeformed configuration]

$$\langle \nabla_\varphi \mathbf{E}; \mathbf{u} \rangle = \frac{1}{2} [\nabla \mathbf{u} + (\nabla \mathbf{u})^T] = \boldsymbol{\epsilon}$$

small-strain tensor
(employed in elasticity theory)



Constitutive relations

Conservation and balance laws

$$J\rho = \rho_0 \quad \forall \mathbf{X} \in B_0 \quad \text{conservation of mass} \quad (1 \text{ equation})$$

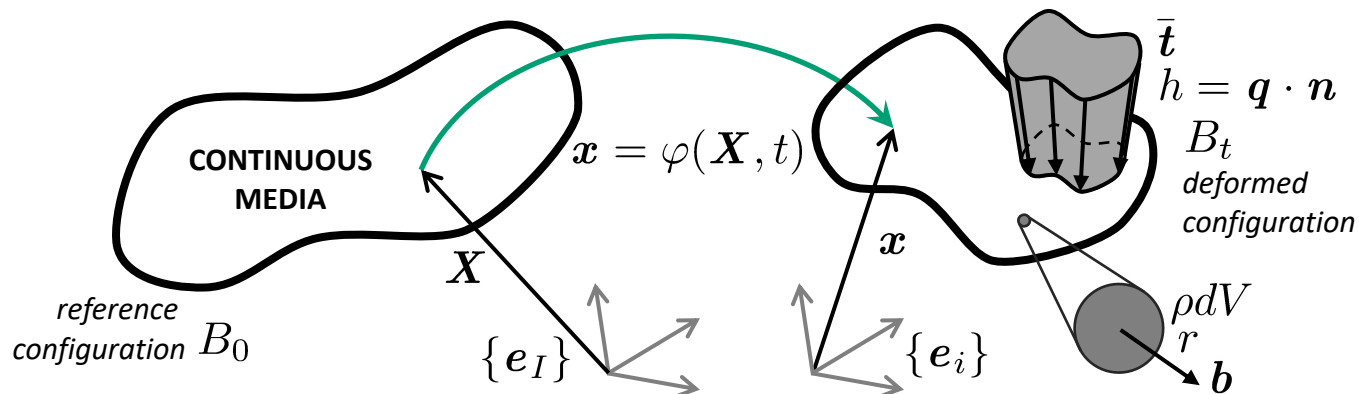
$$\operatorname{div} \boldsymbol{\sigma} + \rho \mathbf{b} = \rho \mathbf{a} \quad \forall \mathbf{x} \in B \quad \text{balance of linear momentum} \quad (3 \text{ equations})$$

$$\boldsymbol{\sigma} = \boldsymbol{\sigma}^T \quad \forall \mathbf{x} \in B \quad \text{balance of angular momentum} \quad (\text{constraint})$$

$$\rho \dot{u} = \boldsymbol{\sigma} : \mathbf{d} + \rho r - \operatorname{div} \mathbf{q} \quad \forall \mathbf{x} \in B \quad \text{conservation of energy} \quad (1 \text{ equation})$$

$$\dot{s} \geq \frac{r}{T} - \frac{1}{\rho} \operatorname{div} \frac{\mathbf{q}}{T} \quad \forall \mathbf{x} \in B \quad \text{Clausius-Duhem inequality} \quad (\text{constraint})$$

$$\rho, \mathbf{x}, \boldsymbol{\sigma}, \mathbf{q}, u, s, T \quad (16 \text{ unknowns})$$



Constitutive relations

Constitutive relations

- Relations that describe the response of the material to mechanical and thermal loading, e.g., $\boldsymbol{\sigma}, \mathbf{q}, W, s$ (11 constitutive equations)

Simple elastic material

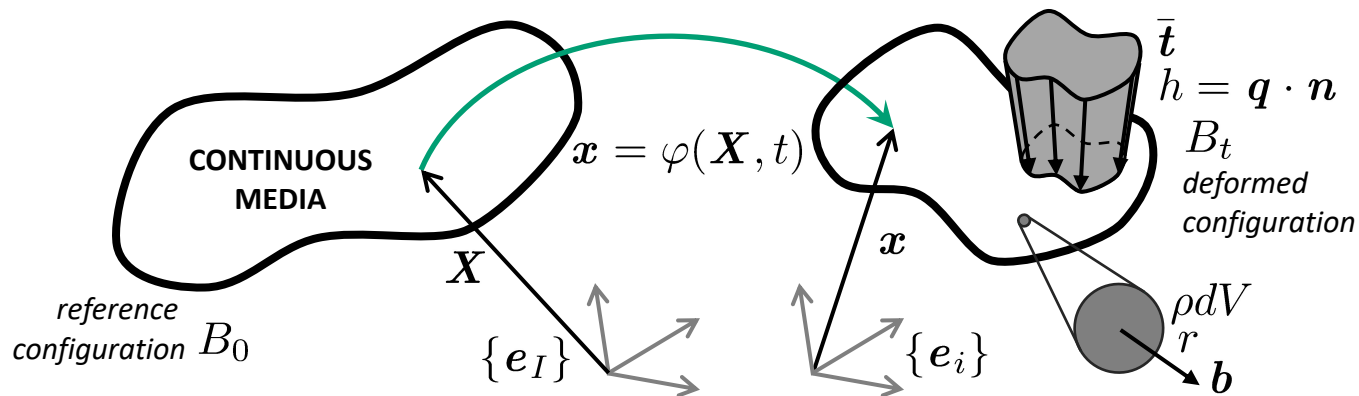
... based on the Helmholtz free energy

$$W = \bar{W}(\mathbf{F}, T)$$

$$\text{entropy } s = \bar{s}(\mathbf{F}, T, \dots)$$

$$\text{heat flux } \mathbf{q} = \bar{\mathbf{q}}(\mathbf{F}, T, \dots)$$

$$\text{Cauchy's stress tensor } \boldsymbol{\sigma} = \bar{\boldsymbol{\sigma}}(\mathbf{F}, T, \dots)$$



Constitutive relations

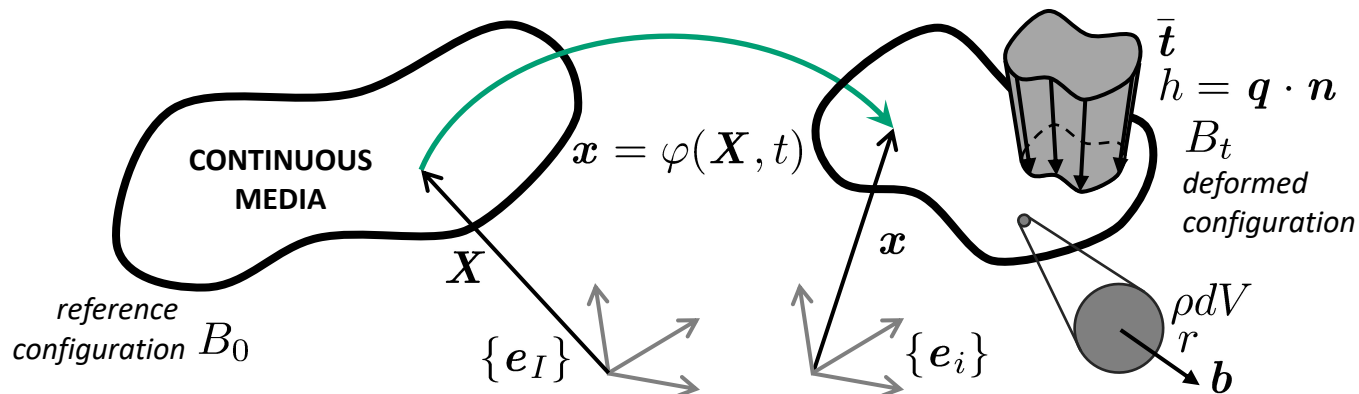
Coleman-Noll procedure

Using conservation of energy, the Clausius-Duhem inequality can be written as

$$\rho T \dot{s}^{\text{int}} = -\rho \left[\frac{1}{\rho_0} \frac{\partial W}{\partial T} + s \right] \dot{T} + \left[\boldsymbol{\sigma} \mathbf{F}^{-T} - \frac{\rho}{\rho_0} \frac{\partial W}{\partial \mathbf{F}} \right] : \dot{\mathbf{F}} - \frac{1}{T} \mathbf{q} \cdot \nabla T \geq 0$$

Coleman and Noll made the argument that this inequality must be satisfied for every admissible process.

The application of the Clausius-Duhem inequality to constitutive equations is known as the Coleman-Noll procedure (1963)



Constitutive relations

Constraints on constitutive relations

- Relations that describe the response of the material to mechanical and thermal loading, e.g., $\boldsymbol{\sigma}, \mathbf{q}, u, T$ (11 constitutive equations)

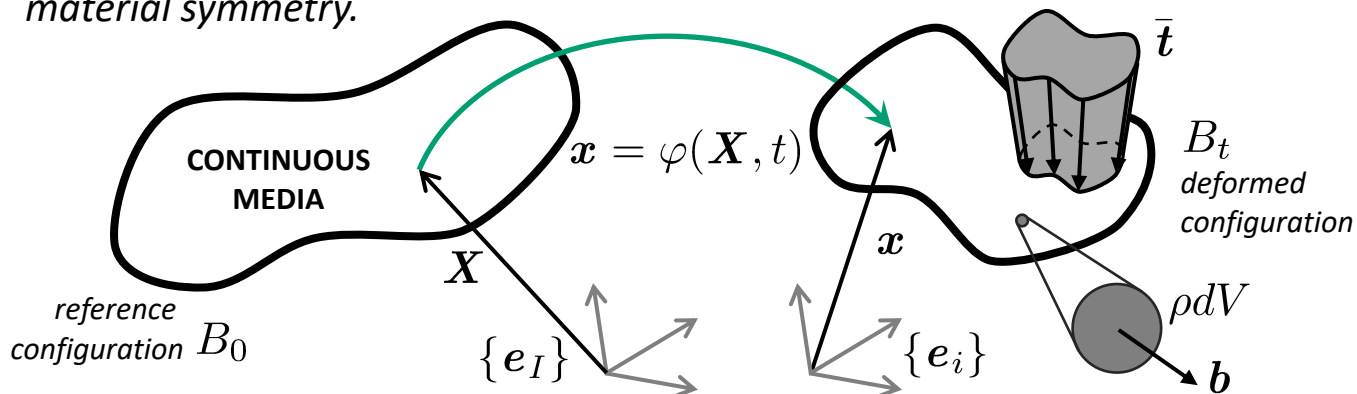
$$\boldsymbol{\sigma} = \boldsymbol{\sigma}^{(e)} + \boldsymbol{\sigma}^{(v)}, \mathbf{q} = \mathbf{0}, \rho_0 u(\mathbf{F}, s), \dot{T} \neq 0 \quad (\text{with } \dot{s} = 0) \quad \text{isentropic processes}$$

$$\boldsymbol{\sigma} = \boldsymbol{\sigma}^{(e)} + \boldsymbol{\sigma}^{(v)}, \mathbf{q} = \mathbf{0}, W(\mathbf{F}, T), \dot{s} \neq 0 \quad (\text{with } \dot{T} = 0) \quad \text{isothermal processes}$$

- Can these constitutive relations be selected arbitrarily? NO!

They must follow the following fundamental principles:

Principle of determinism, principle of local action, second law of thermodynamics restrictions (Clausius-Duhem inequality), principle of material frame indifference, material symmetry.



Constitutive relations

Coleman-Noll procedure – Isothermal processes

- *Isothermal* processes, where the motion and deformation occur at such a low temporal rate that the temperature is uniform and constant.

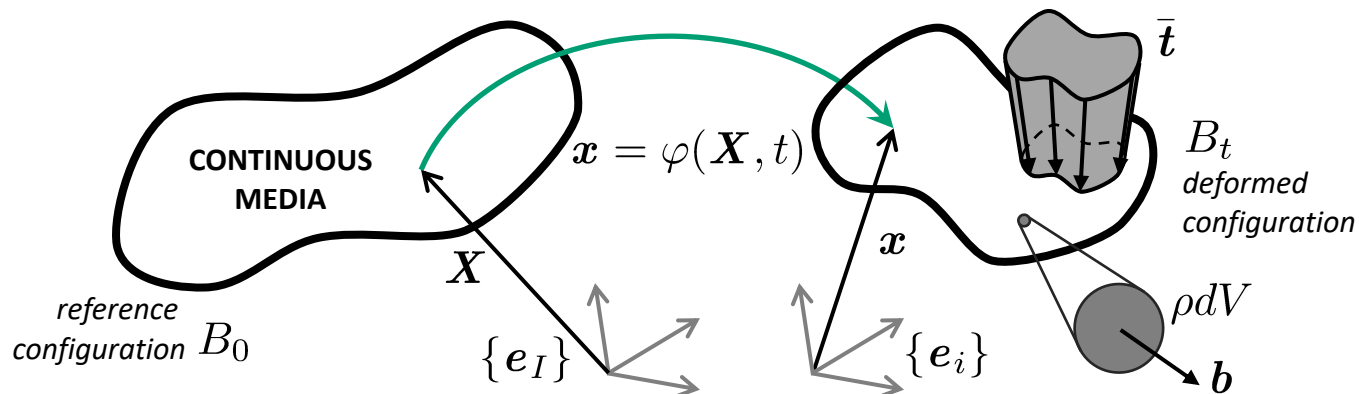
... based on the Helmholtz free energy

$$\dot{T} = 0 \quad \nabla T = 0 \quad \mathbf{q} = -k\nabla T = 0 \quad W = \bar{W}(\mathbf{F}, T)$$

$$s = \bar{s}(\mathbf{F}, T) \equiv -\frac{1}{\rho_0} \frac{\partial W}{\partial T} \quad \dot{s} \neq 0$$

$$\boldsymbol{\sigma}^{(e)} \equiv \frac{1}{J} \frac{\partial W}{\partial \mathbf{F}} \mathbf{F}^T \quad \bar{\boldsymbol{\sigma}}^{(v)}(\mathbf{F}, T, \mathbf{d})$$

If $\boldsymbol{\sigma}^{(v)} = \mathbf{0}$ then the process is reversible, and the material is called **hyperelastic**



Hyperelastic solids - Isotropic

Coleman-Noll procedure + Frame indifference + Isotropy

$$W = W(I_1, I_2, I_3)$$

$$I_1 = \text{tr}(\mathbf{C}) = \text{tr}(\mathbf{B})$$

$$I_2 = \frac{1}{2}[\text{tr}(\mathbf{C})^2 - \text{tr}(\mathbf{C}^2)] = \frac{1}{2}[\text{tr}(\mathbf{B})^2 - \text{tr}(\mathbf{B}^2)]$$

$$I_3 = \det(\mathbf{C}) = \det(\mathbf{B}) = J^2$$

strain energy density function

(a function of the principal invariants of the right/left Cauchy-Green tensor)

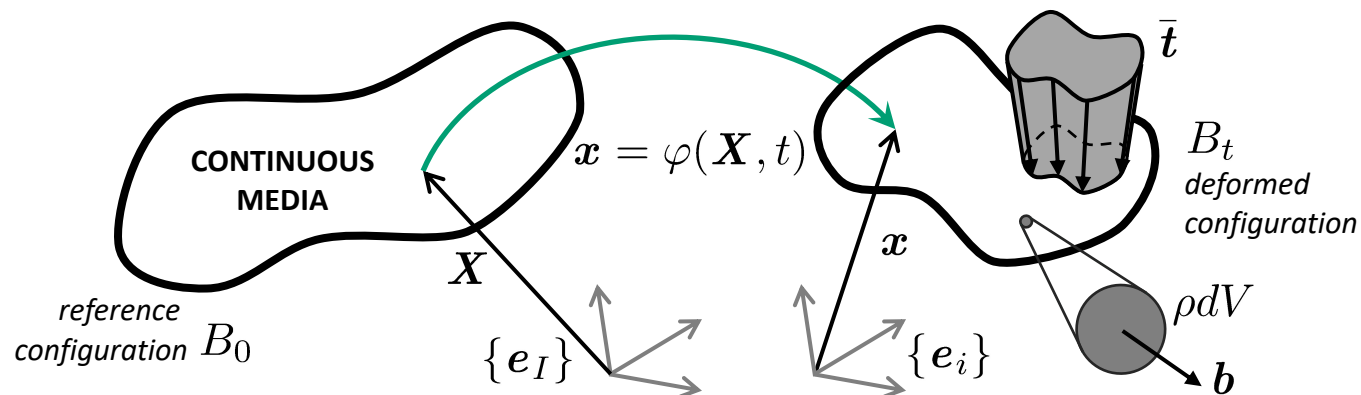
DIY

$$\boldsymbol{\sigma}^{(e)} = \frac{2}{J} \mathbf{F} \frac{\partial \tilde{W}(\mathbf{C}, T)}{\partial \mathbf{C}} \mathbf{F}^T$$

$$\mathbf{S}^{(e)} = 2 \frac{\partial \tilde{W}(\mathbf{C}, T)}{\partial \mathbf{C}}$$

elastic part of the stress tensor

(the viscous part is zero and thus, the process is reversible)



Hyperelastic solids - Isotropic

Coleman-Noll procedure + Frame indifference + Isotropy

$$W = W(I_1, I_2, I_3)$$

$$\mathbf{S} = 2 \left[(W_{,I_1} + I_1 W_{,I_2}) \mathbf{I} - W_{,I_2} \mathbf{C} + I_3 W_{,I_3} \mathbf{C}^{-1} \right]$$

$$\boldsymbol{\sigma} = \frac{2}{I_3^{1/2}} \left[I_3 W_{,I_3} \mathbf{I} + (W_{,I_1} + I_1 W_{,I_2}) \mathbf{B} - W_{,I_2} \mathbf{B}^2 \right]$$

$$\mathbf{S} = 2 \sum_{i=1}^3 \frac{\partial W(I_1, I_2, I_3)}{\partial I_i} \frac{\partial I_i}{\partial \mathbf{C}}$$

$$\frac{\partial I_1}{\partial \mathbf{C}} = \mathbf{I}$$

$$\frac{\partial I_2}{\partial \mathbf{C}} = I_1 \mathbf{I} - \mathbf{C}$$

$$\frac{\partial I_3}{\partial \mathbf{C}} = I_3 \mathbf{C}^{-1}$$

$$W = W(\lambda_1, \lambda_2, \lambda_3)$$

The right and left stretch tensors have the same principal stretches (eigenvalues).

Principal directions (eigenvectors).

$$\boldsymbol{\Lambda}_\alpha^{\mathbf{C}} = \boldsymbol{\Lambda}_\alpha^{\mathbf{U}} = \mathbf{N}_\alpha \quad \alpha = \{1, 2, 3\} \quad \|\mathbf{N}_\alpha\| = 1$$

$$\mathbf{U} \mathbf{N}_\alpha = \lambda_\alpha \mathbf{N}_\alpha \quad \alpha = \{1, 2, 3\}$$

$$\mathbf{C} \mathbf{N}_\alpha = \lambda_\alpha^2 \mathbf{N}_\alpha \quad \alpha = \{1, 2, 3\}$$

$$\frac{\partial \lambda_\alpha}{\partial \mathbf{C}} = \frac{1}{2\lambda_\alpha} \mathbf{N}_\alpha \otimes \mathbf{N}_\alpha \quad \alpha = \{1, 2, 3\}$$

Hyperelastic solids - Isotropic (*and anisotropic*)

Strain energy density function

Isotropic hyperelastic solids.

Blatz-Ko materials (hyperelastic solid)	$W(I_2, I_3) = c_1 \left[\frac{I_2}{I_3} + 2\sqrt{I_3} - 5 \right]$
Neo-Hookean materials (incompressible)	$W(I_1, I_2) = c_1(I_1 - 3) \quad I_3 = 1$
Mooney-Rivlin materials (incompressible)	$W(I_1, I_2) = c_1(I_1 - 3) + c_2(I_2 - 3)$
Ogden model (compressible/incompressible)	$W(\lambda_1, \lambda_2, \lambda_3) = \sum_{p=1}^N \frac{\mu_p}{\alpha_p} (\lambda_1^{\alpha_p} + \lambda_2^{\alpha_p} + \lambda_3^{\alpha_p} - 3)$

Note: incompressibility is an internal constraint (or kinematic constraint) of the material

Question: how is the pressure determined? The undetermined part of the pressure is introduced as a Lagrange multiplier and it is determined from boundary conditions

$$\boldsymbol{\sigma} = \frac{2}{J} \mathbf{F} \frac{\partial \widetilde{W}(\mathbf{C}, T)}{\partial \mathbf{C}} \mathbf{F}^T - c_0 \mathbf{I} \quad \text{with } I_3 = 1 \text{ or } \lambda_3 = 1/\lambda_1 \lambda_2$$

Anisotropic elastic solids. Saint Venant materials. $\widetilde{W}(\mathbf{E}) = \frac{1}{2}(\mathbf{C} : \mathbf{E}) : \mathbf{E}$

$$\mathbf{S} = \frac{\partial \widetilde{W}(\mathbf{E})}{\partial \mathbf{E}} = \mathbb{C} : \mathbf{E} \quad C_{IJKL} = C_{JIKL} = C_{IJLK} = C_{KLIJ}$$

Materials with memory

Internal variable, evolution equation, small strains

- The memory or path-dependency of a material can be represented through an array of **internal variables** (scalars and second-order tensors)

$$\boldsymbol{\sigma}(\boldsymbol{\epsilon}, T, \boldsymbol{\epsilon}^t) \equiv \begin{cases} \boldsymbol{\sigma}(\boldsymbol{\epsilon}, T, \boldsymbol{\xi}) \\ \dot{\xi}_\alpha = g_\alpha(\boldsymbol{\sigma}, T, \boldsymbol{\xi}) \end{cases} \text{ evolution equations}$$

The presence of additional variables in the constitutive laws requires additional constitutive equations, namely **evolution equations**. The hypothesis is that the rate of evolution of the internal variables is also determined from the local state.

Examples:

- **viscoelasticity**, *i.e.*, time- and rate-dependent reversible behavior (the stress–strain relation depends on the loading rate); internal variables, *e.g.*, the inelastic strain.
- **plasticity**, *i.e.*, history-dependent irreversible behavior (the stress–strain relation depends on the loading history); internal variables, *e.g.*, plastic strain, accumulated plastic strain.
- **viscoplasticity**, *i.e.*, history- and time-dependent irreversible behavior.
- **damage**, *i.e.*, irreversible degradation of the elastic stiffness with loading; internal variable is a damage parameter, *e.g.*, a scalar measure.

Materials with memory: Viscoelastic solids

Example of viscoelastic solid: Standard solid model

From equilibrium and compatibility:

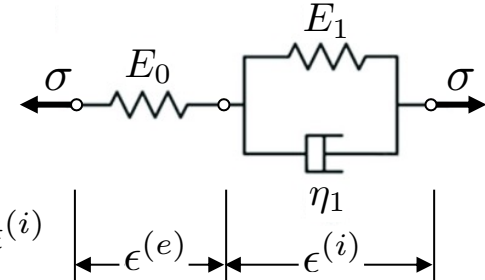
$$\dot{\epsilon}^{(i)} = \frac{1}{\eta_1} \sigma - \frac{E_1}{\eta_1} \epsilon^{(i)}$$

$$\dot{\epsilon}^{(e)} = \frac{E_0}{\eta_1} (\epsilon - \epsilon^{(i)}) - \frac{E_1}{\eta_1} \epsilon^{(i)}$$

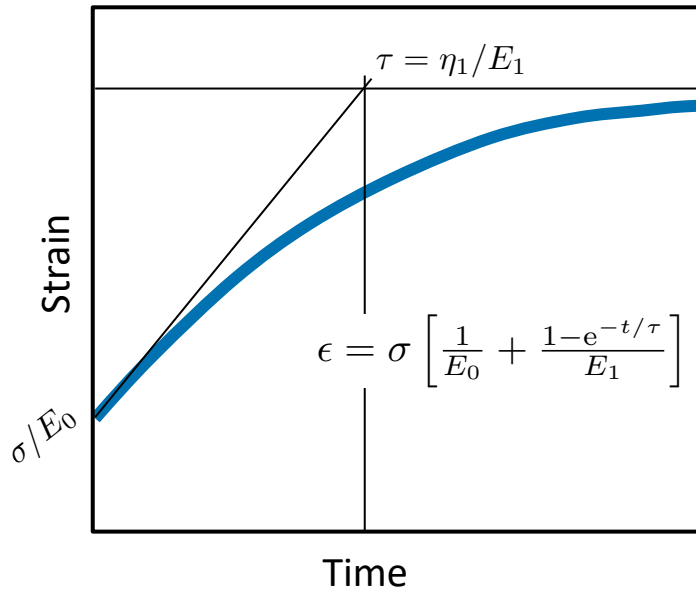
$$\epsilon = \epsilon^{(e)} + \epsilon^{(i)}$$

$$\sigma = E_0 \epsilon^{(e)}$$

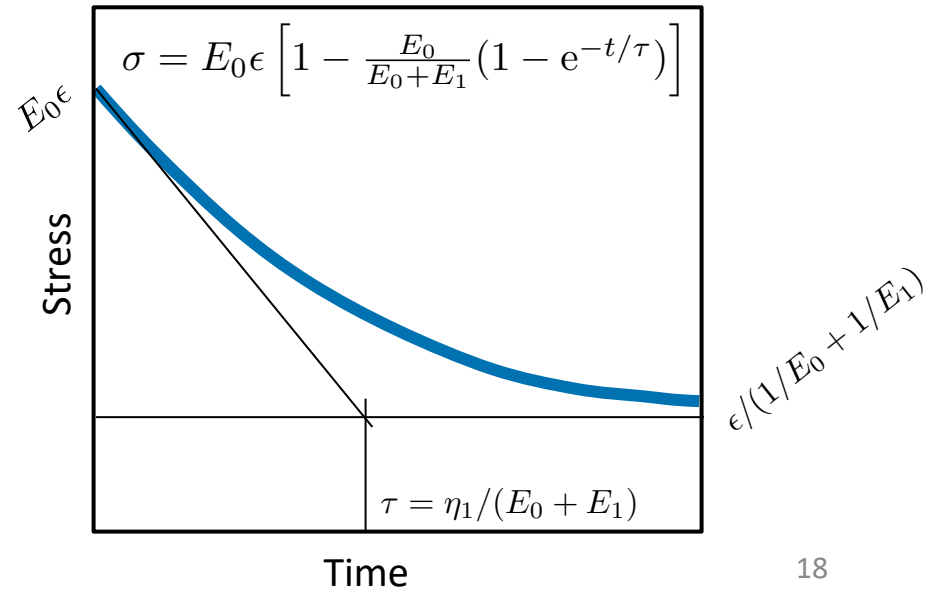
$$\sigma = E_1 \epsilon^{(i)} + \eta_1 \dot{\epsilon}^{(i)}$$



Creep process (at constant stress)



Relaxation process (at constant strain)



Materials with memory: Viscoelastic solids

Internal variable, evolution equation, small strains

- The memory or path-dependency of a material can be represented through an array of **internal variables** (scalars and second-order tensors)

$$\boldsymbol{\sigma}(\boldsymbol{\epsilon}, T, \boldsymbol{\epsilon}^t) \equiv \begin{cases} \boldsymbol{\sigma}(\boldsymbol{\epsilon}, T, \boldsymbol{\xi}) \\ \dot{\boldsymbol{\xi}}_\alpha = g_\alpha(\boldsymbol{\sigma}, T, \boldsymbol{\xi}) \end{cases} \text{ evolution equations}$$

The presence of additional variables in the constitutive laws requires additional constitutive equations, namely **evolution equations**. The hypothesis is that the rate of evolution of the internal variables is also determined from the local state.

- Example: standard solid model

internal variable

$$\xi_1 = \epsilon^{(i)}$$

evolution equation (kinetic equation)

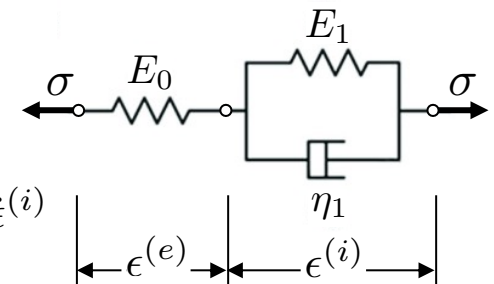
$$\dot{\epsilon}^{(i)} = g_1(\sigma, \xi_1) = \frac{1}{\eta_1} \sigma - \frac{E_1}{\eta_1} \epsilon^{(i)}$$

$$\dot{\epsilon}^{(i)} = g_1(\epsilon, \xi_1) = \frac{E_0}{\eta_1} (\epsilon - \epsilon^{(i)}) - \frac{E_1}{\eta_1} \epsilon^{(i)} \quad (\text{an evolution equation in terms of strain is not always possible})$$

$$\epsilon = \epsilon^{(e)} + \epsilon^{(i)}$$

$$\sigma = E_0 \epsilon^{(e)}$$

$$\sigma = E_1 \epsilon^{(i)} + \eta_1 \dot{\epsilon}^{(i)}$$



Materials with memory

Internal variables and thermodynamics

Using conservation of energy, the Clausius-Duhem inequality can be written as

$$\rho T \dot{s}^{\text{int}} = -\rho \left[\frac{1}{\rho_0} \frac{\partial W}{\partial T} + s \right] \dot{T} + \left[\boldsymbol{\sigma} - \frac{\rho}{\rho_0} \frac{\partial W}{\partial \boldsymbol{\epsilon}} \right] : \dot{\boldsymbol{\epsilon}} - \frac{1}{T} \mathbf{q} \cdot \nabla T \geq 0$$

Coleman and Noll made the argument that this inequality must be satisfied for every admissible process.

Let's now consider path-dependent behavior characterized by a set of internal variables and corresponding evolution equations, that is

$$W = \bar{W}(\boldsymbol{\epsilon}, T, \boldsymbol{\xi}) \quad \dot{\xi}_\alpha = g_\alpha(\boldsymbol{\sigma}, T, \boldsymbol{\xi})$$

- Isentropic/Adiabatic processes $\mathbf{q} = \mathbf{0}$, $W(\boldsymbol{\epsilon}, T, \boldsymbol{\xi})$, $\dot{s} \neq 0$ (with $\dot{T} = 0$)

Assuming $\boldsymbol{\epsilon} = \boldsymbol{\epsilon}^{(e)}(\boldsymbol{\sigma}, T) + \boldsymbol{\epsilon}^{(i)}(\boldsymbol{\xi})$, then there exists $W(\boldsymbol{\epsilon}, T, \boldsymbol{\xi})$

if and only if

$$W(\boldsymbol{\epsilon}, T, \boldsymbol{\xi}) = W^{(e)}(\boldsymbol{\epsilon} - \boldsymbol{\epsilon}^{(i)}(\boldsymbol{\xi}), T) + W^{(i)}(\boldsymbol{\xi}, T)$$

Materials with memory: Viscoelastic solids

Internal variables and thermodynamics

- Example: multiaxial behavior and time integration

$$W(\boldsymbol{\epsilon}, T, \boldsymbol{\xi}) = W^{(e)}(\boldsymbol{\epsilon} - \boldsymbol{\epsilon}^{(i)}(\boldsymbol{\xi}), T) + W^{(i)}(\boldsymbol{\xi}, T)$$

Internal variable $\boldsymbol{\epsilon}^{(i)}(\boldsymbol{\xi}) = \boldsymbol{\xi}$

$$W^{(i)}(\boldsymbol{\xi}, T) = \frac{1}{2} E_1 \boldsymbol{\epsilon}^{(i)} : \boldsymbol{\epsilon}^{(i)}$$

Evolution equation $\dot{\boldsymbol{\epsilon}}^{(i)} = \frac{\boldsymbol{\sigma} - E_1 \boldsymbol{\epsilon}^{(i)}}{\eta_1}$

Time integration $t_n \rightarrow t_{n+1} = t_n + \Delta t$

- Given $\left\{ \boldsymbol{\epsilon}_{n+1}, \boldsymbol{\sigma}_n, \boldsymbol{\epsilon}_n^{(i)} \right\}$

- Update internal variable $\boldsymbol{\epsilon}_{n+1}^{(i)} = \frac{1}{\eta_1/\Delta t} \boldsymbol{\sigma}_n + \boldsymbol{\epsilon}_n^{(i)} \left[1 - \frac{E_1}{\eta_1/\Delta t} \right]$

- Update stress $\boldsymbol{\sigma}_{n+1} = \frac{\rho_{n+1}}{\rho_0} \frac{\partial W_{n+1}^{(e)}}{\partial \boldsymbol{\epsilon}^{(e)}}$

Materials with path-dependency: Plastic solids

Phenomenological observations

- Plasticity manifests when a stress threshold is reached (yield stress)

Yield function

$$\phi(\sigma, \sigma_Y) \leq 0$$

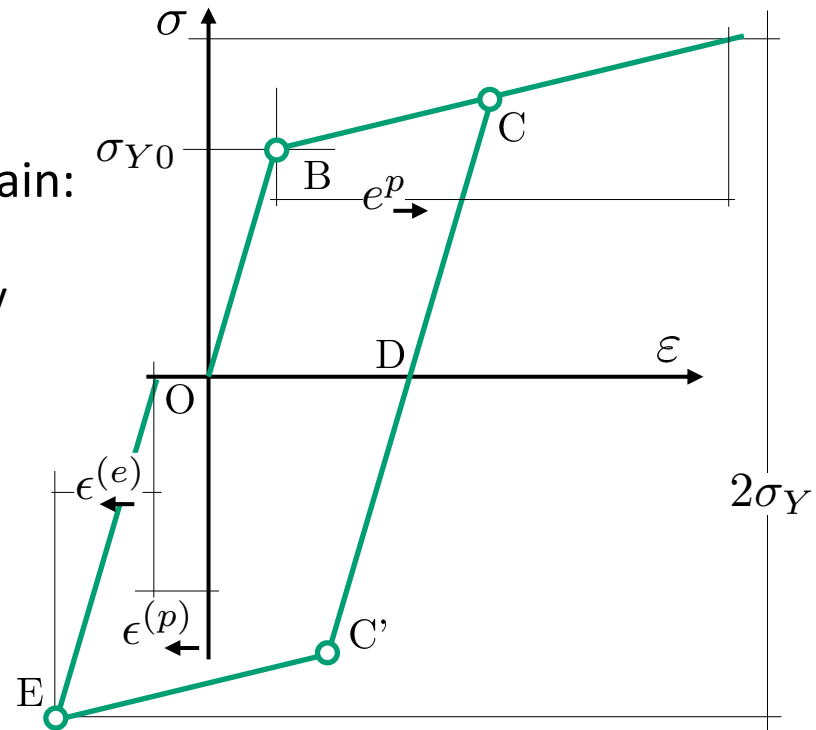
- Additive decomposition of (small) strain:
 - + elastic part is related to the stress
 - + plastic part depends on material history

$$\epsilon = \epsilon^{(e)} + \epsilon^{(p)}$$

- Metals: plastic flow is incompressible (any change in volume is elastic)
- Bauschinger effect $|\sigma_{Y|r}| \leq |\sigma_{Y|f}|$

Strain hardening:

- + isotropic hardening (e^p)
 - + kinematic hardening
 - Metals: Bridgman (c. 1940)
- Yield function is independent of hydrostatic pressure



Isotropic plastic solids

Ideal plastic solids – Small strains

Constitutive law defined by yield function, flow potential and hardening rule

- Plasticity manifests when a stress threshold is reached (yield stress)

Yield function $\phi(\boldsymbol{\sigma}, \xi) \leq 0$

Internal variable (*for isotropic hardening*) $\xi = \int_0^t \|\dot{\boldsymbol{\epsilon}}^{(p)}\| dt$

- Additive decomposition of (small) strain: $\boldsymbol{\epsilon} = \boldsymbol{\epsilon}^{(e)} + \boldsymbol{\epsilon}^{(p)}$

- Consistency condition $\dot{\phi} = 0 \rightarrow$ solve for $\dot{\lambda}$
(*determine $\dot{\lambda}$ under plastic loading; $\dot{\lambda} = 0$, otherwise*)

- Strain energy density: $W(\boldsymbol{\epsilon}, \xi) = W^{(e)}(\boldsymbol{\epsilon} - \boldsymbol{\epsilon}^{(p)}(\xi)) + W^{(p)}(\xi)$

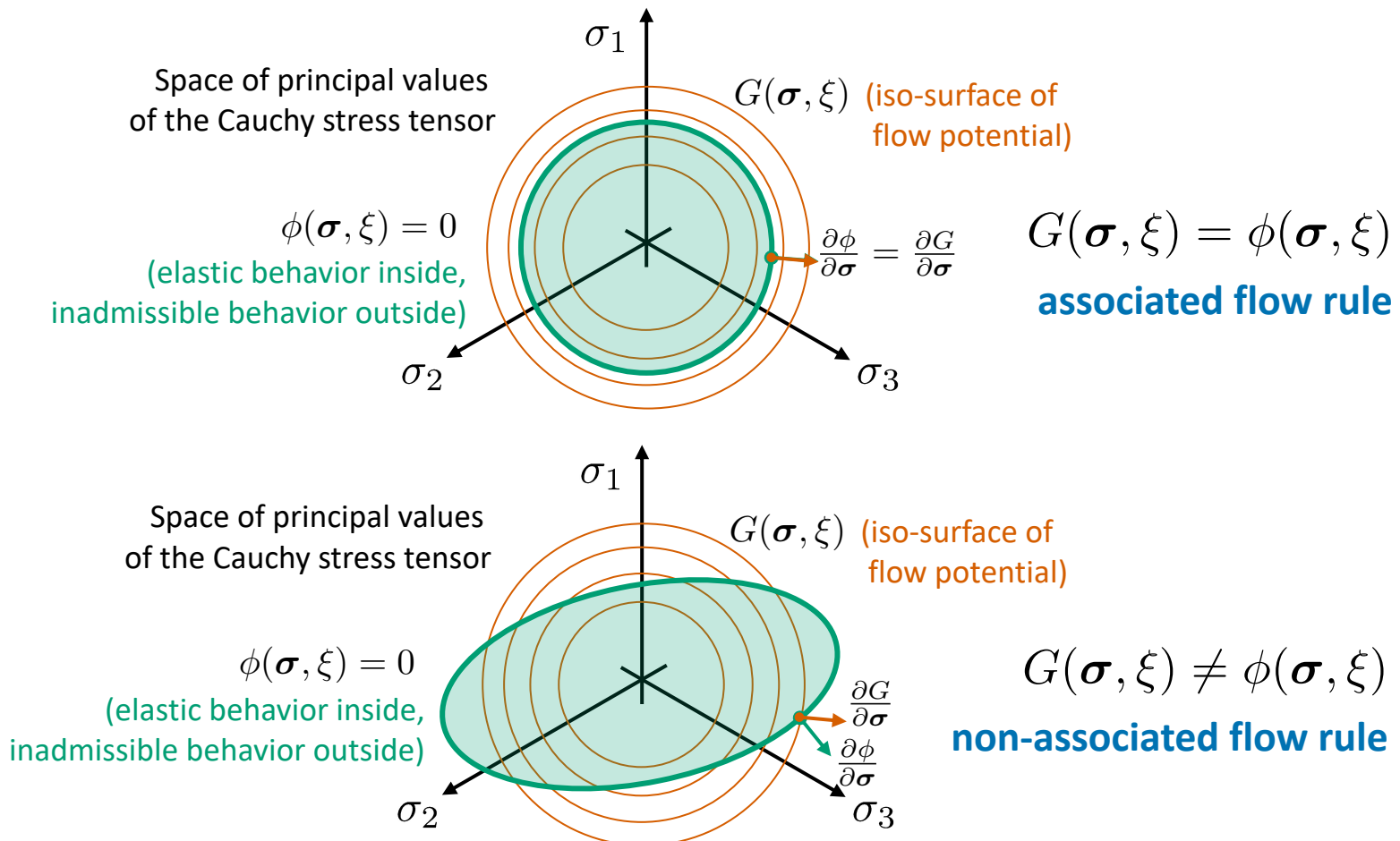
- Evolution equations:
 - Flow rule (flow potential) $\dot{\boldsymbol{\epsilon}}^{(p)} = \dot{\lambda} \mathbf{r}(\boldsymbol{\sigma}, \xi) = \dot{\lambda} \frac{\partial G}{\partial \boldsymbol{\sigma}}$
 - Hardening rule $\dot{\xi} = \dot{\lambda} h(\boldsymbol{\sigma}, \xi)$

- Cauchy stress tensor: $\boldsymbol{\sigma} = \frac{\rho}{\rho_0} \frac{\partial W^{(e)}}{\partial \boldsymbol{\epsilon}^{(e)}} := \mathbf{c}^e : \boldsymbol{\epsilon}^{(e)}$

Isotropic plastic solids

Ideal plastic solids – Small strains

Constitutive law defined by **yield function**, **flow potential** and **hardening rule**



Isotropic plastic solids

Ideal plastic solids – Small strains

Constitutive law defined by yield function, flow potential and hardening rule

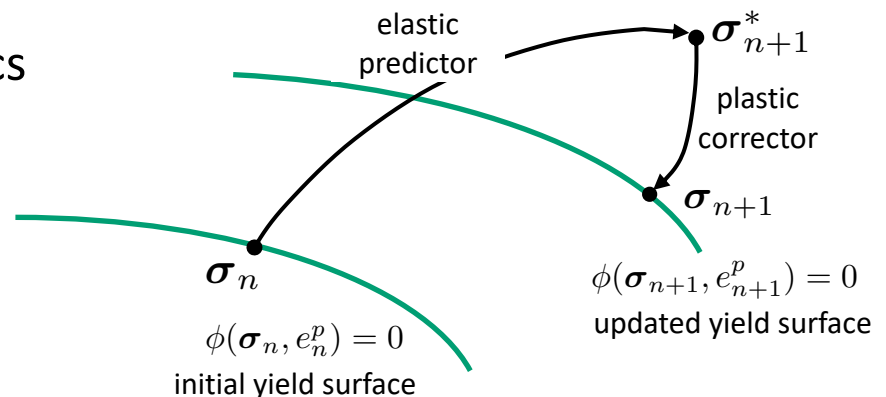
- J_2 or von Mises plasticity
- Drucker-Prager plasticity (associated flow rule)
- Drucker-Prager plasticity (non-associated flow rule)
- Tresca Criterion (or Maximum-shear-stress)
- Mohr-Coulomb plasticity



Q: Which are the chief characteristics of each plastic model?

For example: $\phi(J_2, e^p)$ $\phi(I_1, J_2)$

Q: How is the consistency condition enforced in a time-integration scheme?



Review session for the exam

Study hard!

