Anand Model: Viscoelastoplasticity

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and its Application to Solder Joints

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Constitutive Equations for Hot-Working of Metals

Author: Lallit Anand (1985)

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One of the foundational papers in thermodynamically consistent viscoplasticity modeling—especially significant in the context of metals subjected to large strains and high temperatures.

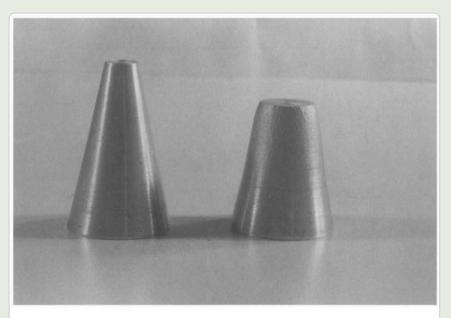


Fig. 25. 1100 aluminum state gradient specimens before and after testing.

International Journal of Masticity, Vol. 1, pp. 213-231, 1965 Printed in the U.S.A.

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CONSTITUTIVE EQUATIONS FOR HOT-WORKING OF METALS

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Massachusetts Institute of Technology

(Communicated by Theoder Lehmann, Ruhr Universität Bochum)

Abstract - Elevated temperature deformation processing - "hot-working," is an important step during the manufacturing of most metal products. Central to any successful analysis of a hot-working process is the use of appropriate rate and temperature-dependent constitutive equations for large, interrupted inelastic deformations, which can faithfully account for strainhardening, the restoration processes of recovery and recrystallization and strain rate and temperature history effects. In this paper we develop a set of phenomenological, internal variable type constitutive equations describing the elevated temperature deformation of metals. We use a scalar and a symmetric, traceless, second-order tensor as internal variables which, in an average sense, represent an isotropic and an anisotropic resistance to plastic flow offered by the internal state of the material. In this theory, we consider small elastic stretches but large plastic deformations (within the limits of texturing) of isotropic materials. Special cases (within the constitutive framework developed here) which should be suitable for analyzing hot-working processes are indicated.

I. INTRODUCTION

Hot-working is an important processing step during the manufacture of approximately more than eighty-five percent of all metal products. The main features of hot-working are that metals are deformed into the desired shapes at temperatures in the range of −0.5 through ~0.9 θ_m, where θ_m is the melting temperature in degrees Kelvin, and at strain rates in the range of -10-4 through -103/sec. It is to be noted that most hotworking processes are more than mere shape-making operations; an important goal of hot-working is to subject the workpiece to appropriate thermo-mechanical processing histories which will produce microstructures that optimize the mechanical properties of the product.

The major quantities of metals and alloys are hot-worked under interrupted nonisothermal conditions. The principles of the physical metallurgy of such deformation processing are now well recognized, e.g., Jonas et al. [1969], Sellars & McG Tegart [1972], McQueen & Jonas [1975], and Sellars [1978]. During a deformation pass, the stress is found to be a strong function of the strain rate, temperature, and the defect and microstructural state of the material. The strain-hardening produced by the deformation tends to be counteracted by dynamic recovery processes. These recovery processes result in a rearrangement and annihilation of dislocations in such a manner that as the strain in a pass increases, the dislocations tend to arrange themselves into subgrain walls. In some metals and alloys (especially those with a high stacking fault energy, e.g., Al, α-Fe and other ferritic alloys) dynamic recovery can balance strain-hardening and an apparent steady state stress level can be achieved and maintained to large strains before fracture occurs. In other metals and alloys in which recovery is less rapid (especially those metals with low stacking fault energies, e.g., Ni, γ-Fe and other austenitic

Case Study: Wang (2

G. Z. Wang

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Applying Anand Model to Represent the Viscoplastic **Deformation Behavior of Solder** Alloys

A unified viscoplastic constitutive law, the Annual model, was applied to represent the industric deformation behavior for solders used in electronic packaging. The material parameters of the constitutive relations for 62x8/69924g, 605x6099, 95.58x3.5g, and 95.75492.28s tolders were determined from separated constitutive relations and experimental results. The achieved unified Annual model for solders were texted for constitution.

Introduction

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Introduction

Int

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SEPTEMBER 2001, Vol. 123 / 247

Source: Wang, C. H. (2001). "A Unified Creep-Plasticity Model for Solder Alloys."

DOI: 10.1115/1.1371781

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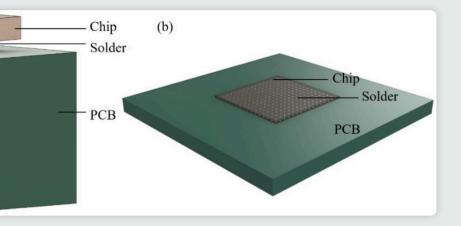
Why Wang's Paper Matters

and's unified viscoplastic framework to model solder behavior.

odel can be reduced and fitted from experiments.

he theory into engineering-scale implementation.

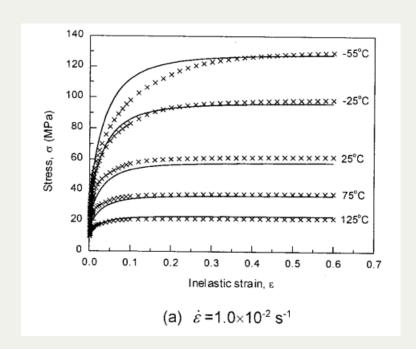
lder joints in microelectronic packages (chip on PCB, soldered as).



Observed Behavior

- Top Graph (a): $\dot{arepsilon}=10^{-2}\,\mathrm{s}^{-1}$
- High strain rate → higher stress
- Recovery negligible → pronounced hardening
- Bottom Graph (b): $\dot{arepsilon}=10^{-4}\,\mathrm{s}^{-1}$
- Lower strain rate → lower stress at same strain
- · Recovery and creep effects more significant

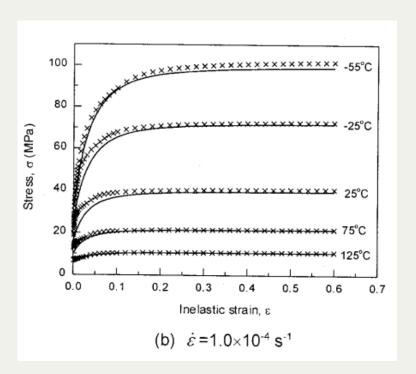
Model Accuracy: Lines = model prediction, X = experimental data



Key Insights from Wang (2001)

- "At lower strain rates, recovery dominates... the stress levels off early."
- "At high strain rates, hardening dominates, and the stress grows continuously."

Anand's model smoothly captures strain-rate and temperature dependence of solder materials.



Flow Rule (Plastic Strain Rate)

$$\dot{arepsilon}^p = A \exp igg(-rac{Q}{RT} igg) igg[\sinh igg(rac{j\sigma}{s} igg) igg]^{1/m}$$

- Plastic strain rate increases with stress and temperature.
- No explicit yield surface; flow occurs at all nonzero stresses.

Deformation Resistance Saturation s^*

$$oldsymbol{s}^* = \hat{s} \left(rac{\dot{arepsilon}^p}{A} ext{exp} \left(rac{Q}{RT}
ight)
ight)^n$$

- Defines the steady-state value that s evolves toward.
- Depends on strain rate and temperature.

and-Type Viscoplastic Model

Evolution of Deformation Resistance *s*

$$\dot{s} = h_0 \Big| 1 - rac{s}{s^*} \Big|^a \operatorname{sign} \left(1 - rac{s}{s^*}
ight) \dot{arepsilon}^p$$

- Describes dynamic hardening and softening of the material.
- s evolves depending on proximity to s^{*} and flow activity.

Note: Constants $A, Q, m, j, h_0, \hat{s}, n, a$ are material-specific and fitted to experimental creep/strain rate data.

Image Reference

Values are from correspond to 60Sn40Pb solder parameters used in Anand's model:

- S_0 : Initial deformation resistance
- Q/R: Activation energy over gas constant
- A: Pre-exponential factor for flow rate
- ξ: Multiplier of stress inside sinh
- *m*: Strain rate sensitivity of stress
- *h*₀: Hardening/softening constant
- \$\hat{s}\$: Coefficient for saturation stress
- n: Strain rate sensitivity of saturation
- a: Strain rate sensitivity of hardening or softening

Constants for 60Sn40Pb

Numerical Values

•
$$S_0 = 5.633 \times 10^7 \, {
m Pa}$$

•
$$Q/R = 10830 \text{ K}$$

•
$$A = 1.49 \times 10^7 \text{ s}^{-1}$$

•
$$\xi = 11$$

•
$$m = 0.303$$

•
$$h_0 = 2.6408 imes 10^9 \; \mathsf{Pa}$$

•
$$\hat{s} = 8.042 \times 10^7 \, \text{Pa}$$

•
$$n = 0.0231$$

•
$$a = 1.34$$

These constants match Wang's paper for modeling 60Sn40Pb viscoplasticity.

Forward Euler Explicit time in

Initialization

- Material constants: $A,Q/R,j,m,h_0,\hat{s},n,a,E$
- Strain rate: $\dot{\varepsilon}$
- Temperature set: $\{T_i\}$
- Set: $\varepsilon^p(0) = 0$, $s(0) = \hat{s}$

Time Evolution Loop

- 1. $\varepsilon_{\mathrm{total}}(t) = \dot{\varepsilon} t$
- 2. $\sigma_{\mathrm{trial}} = E(\varepsilon_{\mathrm{total}} \varepsilon^p)$
- 3. Compute $x = \frac{j\sigma}{s}$
- 4. Approximate $\sinh(x)$ (linearize if $|x|\ll 1$)
- 5. $\dot{\varepsilon}^p = Ae^{-Q/RT}(\sinh(x))^{1/m}$

ntegration scheme Pseudocode

Plastic Flow & Resistance Evolution

6.
$$s^* = \hat{s} \left(rac{\dot{arepsilon}^p}{A} e^{Q/RT}
ight)^n$$

7.
$$\dot{s} = h_0 \left| 1 - \frac{s}{s^*} \right|^a \text{sign} \left(1 - \frac{s}{s^*} \right) \dot{\varepsilon}^p$$

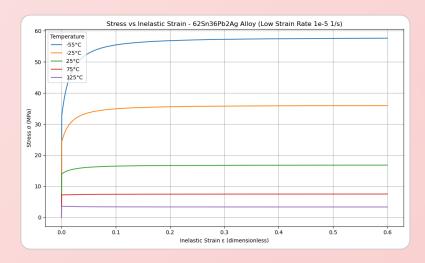
8. Update:
$$\varepsilon^p(t+\Delta t) = \varepsilon^p(t) + \dot{\varepsilon}^p \Delta t$$

9. Update:
$$s(t+\Delta t)=s(t)+\dot{s}\Delta t$$

10. Record $(\varepsilon_{\text{total}}, \sigma_{\text{trial}})$

Termination

- Stop when $arepsilon_{ ext{total}} \geq arepsilon_{ ext{max}}$
- Plot σ vs ε for all T_i

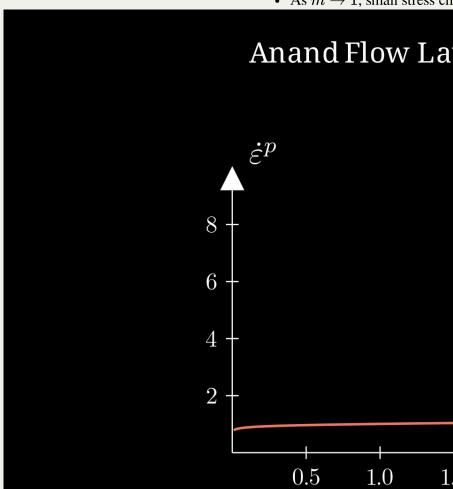


```
import numpy as np
import matplotlib.pyplot as plt
from scipy.integrate import solve ivp
# Material constants for 62Sn36Pb2Ag solder alloy
A = 2.24e8 # 1/s
Q_R = 11200
                 # K
j = 13
                 # dimensionless
m = 0.21
                 # dimensionless
m = 0.21
h0 = 1.62e10  # Pa
s0 = 8.47e7  # Pa
s0 = 8.47e7
s hat = 8.47e7 # Pa
= 0.0277 # dimensionless
a = 1.7 # dimensionless
E = 5.2e10 # Pa (Elastic modulus)
# Temperatures in Kelvin
T_C = [-55, -25, 25, 75, 125]
T_list = [T + 273.15 for T in T_C]
# Simulation parameters
strain rate = le-5 # 1/s
eps total max = 0.6
t max = eps total max / strain rate
time steps = 100
t_eval = np.linspace(0, t_max, time_steps)
# Define the ODE system
def system(t, y, T):
    ep_p, s = y
    eps_total = strain_rate * t
    sigma_trial = E * (eps_total - ep_p)
x = j * sigma_trial / s
    if np.abs(x) < 0.01:
        sinh_x = x
    else:
         sinh_x = np.sinh(np.clip(x, -30, 30))
    sinh x = np.maximum(sinh x, 1e-12)
    dep_p = A * np.exp(-Q_R / T) * sinh_x**(1/m)
    s_star = s_hat * (dep_p / A * np.exp(Q_R / T))**n
    ds = h0 * np.abs(1 - s/s_star)**a * np.sign(1 - s/s_star) * dep_p
    return [dep_p, ds]
# Plotting
```

me for Anand Model

Strain rate sensit

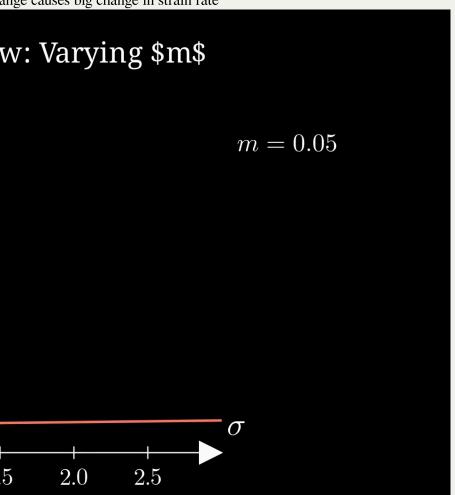
- As m o 0, rate insensitive
- As m o 1, small stress ch



civity of stress m

(yield)

ange causes big change in strain rate



Tensorial Flow Rule (directional form)

$$\mathbf{D}^p = \dot{\epsilon}^p \left(rac{3}{2} rac{\mathbf{T}'}{ar{\sigma}}
ight)$$

Equivalent Stress Definition

$$ar{\sigma} = \sqrt{rac{3}{2} \mathbf{T}' : \mathbf{T}'}$$

- Direction given by \mathbf{T}' .
- Magnitude determined by hyperbolic
- $\bar{ au}$ represents the effective shear stre
- $\bar{\sigma} = \sqrt{\frac{3}{2} \mathbf{T}' : \mathbf{T}'}$ is the von Mises Eq

Summary:

• Full flow = direction × magnitude.

rule

Plastic Strain Rate (magnitude form)

$$\dot{\epsilon}^p = A \exp igg(-rac{Q}{R heta} igg) igg[\sinh igg(\xi rac{ar{\sigma}}{s} igg) igg]^{1/m}$$

Full Flow Rule with Hyperbolic Sine

$$\mathbf{D}^p = A \exp\!\left(-rac{Q}{R heta}
ight)\!\left[\sinh\!\left(\xirac{ar{\sigma}}{s}
ight)
ight]^{1/m} \left(rac{3}{2}rac{\mathbf{T}'}{ar{\sigma}}
ight),$$

$$ar{m{ au}}=\dot{m{\gamma}}^p\left(rac{\mathbf{ ilde{T}}'}{2ar{ au}}
ight),\quad ar{ au}=\left\{rac{1}{2}\mathrm{tr}(\mathbf{ ilde{T}}'^2)
ight\}^{1/2}.$$

sine based on $ar{\sigma}/s$.

ss computed from deviatoric stress.

uivalent stress, but is formally defined without yield point

Stress Evolution Equation (Rate form of Hooke's Law)

$$\overset{
abla}{\mathbf{T}} = \mathbb{L}\left[\mathbf{D} - \mathbf{D}^p
ight] - \mathbf{\Pi}\dot{ heta}$$

(rate-form Hooke's law for finite deformation plasticity, with frame-indifference enforced through the Jaumann rate.)

Jaumann Rate Definition

$$\overset{ riangledown}{\mathbf{T}}=\dot{\mathbf{T}}-\mathbf{W}\mathbf{T}+\mathbf{T}\mathbf{W}$$

Summary:

- Stress rate follows Jaun
- Elastic response govern
- Thermal expansion intro

ion for the Stress

Material Tensors and Operators

- $\mathbb{L}=2\mu \mathbf{I}+\left(\kappa-rac{2}{3}\mu
 ight)\mathbf{1}\otimes\mathbf{1}$ isotropic elasticity tensor
- LD represents how instantaneous strain rates generate stresses according to the elastic material's stiffness properties.
- $\mu = \mu(\theta)$, $\kappa = \kappa(\theta)$ temperature-dependent moduli
- $\Pi = (3\alpha\kappa)\mathbf{1}$ stress-temperature coupling
- $\alpha = \alpha(\theta)$ thermal expansion coefficient
- $\mathbf{D} = \operatorname{sym}(\nabla \mathbf{v})$ stretching tensor
- $\mathbf{W} = \operatorname{skew}(\nabla \mathbf{v})$ spin tensor
- I = fourth-order identity tensor
- 1 = second-order identity tensor

nann derivative to ensure frame indifference.

ed by isotropic fourth-order tensor $\mathbb{L}.$

duces additional stress through ${f \Pi}\dot{ heta}$.

Stress Evolution and Thermal Effects

In the stress evolution equation,

$$\overset{
abla}{\mathbf{T}} = \mathbb{L}\left[\mathbf{D} - \mathbf{D}^p
ight] - \mathbf{\Pi}\dot{ heta},$$

the term $\Pi\dot{\theta}$ represents the stress change that would occur due to pure thermal expansion alone, without any mechanical loading.

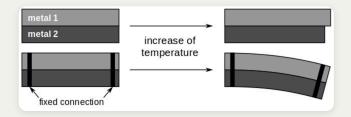
Summary:

- Thermal expansion induces st
 Subtracting Πθ ensures only r
- Subtracting $\mathbf{1}\mathbf{1}\theta$ ensures only
- This keeps the constitutive mo

nd Thermal Effects

Why Subtract the Thermal Term?

- Thermal expansion creates strain even without external forces.
- Without subtracting $\Pi \dot{\theta}$, the model would falsely attribute thermal strain as mechanical stress.
- Subtracting isolates the true mechanical response from thermal effects.



rain without force.

nechanical strains generate stresses.

del physically accurate during heating and cooling.

Context for the Relaxed Configuration

- The relaxed configuration represents the material after removing plastic deformations but before applying new elastic deformations.
- It is introduced to separate permanent plastic effects from recoverable elastic effects.
- All thermodynamic potentials, internal variables, and evolution laws are defined relative to this frame.
- The relaxed state provides a clean, natural reference for measuring elastic strain E^e and computing dissipation.

Sum

• The relaxed configuration isolates elastic responses plastic evolution laws.

iate) Configuration

What Happens in the Relaxed Configuration?

- The elastic deformation gradient F^e is measured from the relaxed state to the current deformed state.
- Elastic strain measures like C^e and E^e are defined in this configuration.
- The Kirchhoff stress $\widetilde{\mathbf{T}}$ is naturally associated with the relaxed volume.
- Plastic flow is accounted for separately through the plastic velocity gradient L^p.

mary:

cleanly, enabling proper definition of thermodynamics and

Kinematics in the Relaxed Configuration

· Elastic deformation gradient:

$$F = F^e F^p \quad \Rightarrow \quad F^e = F F^{p-1}$$

• Elastic right Cauchy-Green tensor:

$$C^e = F^{eT}F^e$$

· Elastic Green-Lagrange strain tensor:

$$E^e = \frac{1}{2}(C^e - I)$$

Sum

- Elastic kinematics and stress measures are formulated plastic and elastic contributions.
 Stress Power Split allows Apand to cleanly isolate plant.
- Stress Power Split allows Anand to cleanly isolate pla
- Green-Lagrange strain tensor E^e is used because it relaxed configuration
- The right Cauchy-Green tensor $C^e = F^{e^T}F^e$ is required deformation gradient F^e without referencing spatial of

n Constituative Laws

Stress and Power Quantities

• Kirchhoff stress (weighted Cauchy stress):

$$\widetilde{\mathbf{T}} = (\det F)\mathbf{T}$$

Stress power split:

$$\dot{\omega} = \dot{\omega}^e + \dot{\omega}^p$$

$$\dot{\omega}^e = \widetilde{f T} : \dot{E}^e \quad , \quad \dot{\omega}^p = (C^e \widetilde{f T}) : {f L}^p$$

mary:

ed relative to the relaxed configuration, cleanly separating

astic dissipation from elastic storage.

symmetrically captures nonlinear elastic strain relative to the

red as an intermediate to compute E^e from the elastic oordinates

Thermodynamic Separation

1. Start with Total Dissipation:

$$\mathcal{D}=\dot{\omega}-\dot{\psi}>0$$

where $\dot{\omega}=\mathbf{\widehat{T}}:\mathbf{\dot{E}}^{e}+(\mathbf{C}^{e}\mathbf{\widehat{T}}):\mathbf{L}^{p}$

2. Split Stress Power:

$$\dot{\omega}=\dot{\omega}^e+\dot{\omega}^p$$

with:

- $oldsymbol{\dot{\omega}}^e = \widehat{f T} : \dot{f E}^e$
- $oldsymbol{\dot{\omega}}^p = (\mathbf{C}^e \widehat{\mathbf{T}}) : \mathbf{L}^p$

3. Group Terms with $\dot{\psi}$:

$$(\dot{\omega}^e - \dot{\psi}) + \dot{\omega}^p \geq 0$$

4. Apply Elastic Energy Consistency:

$$\dot{\omega}^e - \dot{\psi} = 0 \quad \Rightarrow \quad \dot{\omega}^p \geq 0$$

Key Physical Insights

- Elastic deformations are recoverable and do not cause entropy production.
- All dissipation stems from the plastic flow: $\dot{\omega}^p$.
- Plastic work increases entropy and governs viscoplastic evolution.

Summary:

The stress power split ensures that the second law is satisfied by assigning dissipation solely to irreversible processes.

Framework in the Reference Configuration

- The free energy ψ is defined relative to the reference configuration.
- State variables like $E^e, \theta, \bar{g}, \bar{\mathbf{B}}, s$ are used as arguments of ψ .
- Stress is expressed using the second Piola–Kirchhoff tensor S.
- Dissipation inequality, stress—strain relations, and evolution laws are all written in reference variables.
- Mass density ρ_0 from the reference configuration normalizes all terms.

Sum

In the reference configuration, all energy storage, strewith reference-frame quantities for consistency and configuration.

onfiguration

Key Equations in the Reference Frame

• Free energy:

$$\boxed{\psi = \psi(E^e, heta, ar{g}, ar{\mathbf{B}}, s)}$$

· Dissipation inequality:

$$oxed{\dot{\psi} + \eta \dot{ heta} -
ho_0^{-1} \mathbf{S} : \dot{E} + (
ho_0 heta)^{-1} \mathbf{q}_0 \cdot \mathbf{g}_0 \leq 0}$$

· Constitutive relation:

$$\mathbf{S}=
ho_0rac{\partial \psi}{\partial E^e}$$

mary:

ess updates, and internal variable evolution are formulated bjectivity.

Thermodynamic Quantities

• Free energy density:

$$\boxed{\psi = \epsilon - heta \eta}$$

Reduced dissipation inequality:

$$oxed{\dot{\psi} + \eta \dot{ heta} -
ho^{-1} \mathbf{T} : \mathbf{L} + (
ho heta)^{-1} \mathbf{q} \cdot \mathbf{g} \leq 0}$$

· State variables:

$$\{E^e, heta, ar{g}, ar{\mathbf{B}}, s\}$$

with E^e as elastic strain and s as internal resistance.

- Free energy and dissipation
- Stress power naturally split
- Summary: Kirchhoff stress simplifies s

lynamics

Stress Power and Kirchhoff Stress

• Stress power per relaxed volume:

$$\left[\dot{\omega} = \left(rac{
ho_0}{
ho}
ight) \mathbf{T}: \mathbf{L}
ight]$$

• Weighted Cauchy (Kirchhoff) stress:

$$oxed{\widetilde{\mathbf{T}} = (\det F)\mathbf{T}} \quad ext{or} \quad oxed{\widetilde{\mathbf{T}} = \left(rac{
ho_0}{
ho}
ight)\mathbf{T}}$$

Decomposition of stress power:

$$\dot{\omega}=\dot{\omega}^e+\dot{\omega}^p$$

$$\dot{\omega}^e = \widetilde{\mathbf{T}} : \dot{E}^e, \quad \dot{\omega}^p = (C^e \widetilde{\mathbf{T}}) : \mathbf{L}^p$$

n govern thermodynamic consistency.

s into elastic and plastic parts.

tress evolution accounting for volume changes.