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AN EXTENDED MECHANICAL THRESHOLD STRESS PLASTICITY MODEL: MODELING 6061-T6 ALUMINUM ALLOY

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The mechanical threshold stress plasticity model of Follansbee and Kocks was designed to predict the flow stress of metals and alloys in the regime where thermally activated mechanisms are dominant and high temperature diffusion effects are negligible. In this paper we present a model that extends the original mechanical threshold stress to the high strain-rate regime (strain rates higher than 10^4 s^{-1}) and attempts to allow for high temperature effects. We use a phonon drag model for moderate strain rates and an overdriven shock model for extremely high strain rates. A temperature dependent model for the evolution of dislocation density is also presented. In addition, we present a thermodynamically-based model for the evolution of temperature with plastic strain. Parameters for 6061-T6 aluminum alloy are determined and compared with experimental data. The strain-rate dependence of the flow stress of 6061-T6 aluminum is found to be in excellent agreement with experimental data. The amount of thermal softening is underestimated at high temperatures (greater than 500 K) but still is an improvement over the original model. We also find that the pressure dependence of the shear modulus does not completely explain the pressure dependence of the flow stress of 6061-T6 aluminum alloy.

1. Introduction

With the advent of powerful computational machines and algorithms, it is now possible to simulate processes such as the accidental impact of vehicles, high-speed machining, high- and hyper-velocity impact and penetration, and explosive forming. These processes typically involve plastic deformations at high strain rates (10^2 s^{-1} to 10^9 s^{-1}) and often temperatures greater than $0.6 T_m$ (T_m is the melting temperature of the material). Adiabatic shear banding often accompanies the process and provides zones where fractures nucleate. A crucial component in the numerical modeling of plastic deformations under these conditions is a model that describes the evolution of the flow stress under a large range of strain-rates and temperatures.

The mechanical threshold stress (MTS) flow stress model [Follansbee and Kocks 1988; Goto et al. 2000; Kocks 2001] was originally developed with two assumptions:

- (1) Thermally activated dislocation motion was dominant and viscous drag effects on dislocation motion were small. This assumption essentially restricts the model to strain rates of 10^4 s^{-1} and less.
- (2) High temperature diffusion effects (such as solute diffusion from inside the grains to grain boundaries) were absent. This assumption limits the range of applicability of the model to temperatures

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less than around $0.6 T_m$ (depending on the material). For 6061-T6 aluminum alloy this temperature is approximately 450–500 K.

The aluminum alloy 6061 is composed of 97.5% aluminum, 1.0% magnesium, 0.6% silicon, 0.34% iron, 0.3% copper, 0.2% chromium, and around 0.04% zinc and titanium by weight. The T6 hardening treatment varies by manufacturer but usually involves a solution treatment at 810 K followed by quenching in water and then aging for a few hours at around 450 K. The solution treatment leads to a homogeneous supersaturated solid solution. The quenching step is performed to take the supersaturated solution to a two-phase region of the phase diagram. In the aging step, the magnesium silicide (Mg_2Si) phase is precipitated in such a way that the precipitate is evenly distributed inside the grains. There is also some precipitation of $AlFeSi$ at the grain boundaries. The precipitates inside the grains impede dislocation motion and make the alloy harder. The lightness, strength, weldability, and corrosion resistance of the 6061-T6 alloy makes it ideal for heavy duty structures in marine applications (among other uses). However, at about 550 K, the alloying Mg_2Si precipitates start to diffuse toward the grain boundaries and accumulate at the grain boundaries. As a result, 6061-T6 aluminum softens considerably at high temperatures. More details about the processes involved can be found in [Epler 2002].

Simulations of high velocity sphere-cylinder impact experiments using a MTS model for an aluminum alloy containing 1% magnesium [Puchi-Cabrera et al. 2001] showed that the model underestimated the flow stress at strain rates greater than $10^5 s^{-1}$. Also, high temperature Taylor impact simulations using the same model showed that the flow stress was significantly overestimated for temperatures greater than 450 K [Banerjee 2005b].

In this paper, we extend the MTS model so that it can be used for strain rates greater than $10^4 s^{-1}$ and for temperatures greater than 450 K. To extend the model to the regime of high strain rates, we incorporate a model for viscous drag controlled dislocation motion (which becomes dominant at strain rates greater than $10^4 s^{-1}$). In addition, we incorporate a model for the overdriven shock regime (strain-rates greater than $10^9 s^{-1}$). To account for the effect of thermally assisted diffusion of solutes, we incorporate an empirical modification of the original model for thermally activated mechanisms.

The evolution of temperature with plastic strain can be quite important in some situations (such as the development of adiabatic shear bands). We derive an expression for the Taylor–Quinney coefficient from thermodynamic principles and postulate a temperature dependent model for the evolution of this quantity.

The parameters of the extended MTS model are then fit using experimental stress-strain data for 6061-T6 aluminum from the literature wherever available. The flow stress estimated from the model is then compared with experimental data. Next, predictions of the model for a range of strain rates, temperatures, and levels of plastic strain are discussed. Finally, some concluding remarks are given and potential areas of future work are discussed.

2. The extended mechanical threshold stress model

The original mechanical threshold stress (MTS) model [Follansbee and Kocks 1988; Goto et al. 2000] for the flow stress (σ_y) can be written as

$$\sigma_y(\sigma_e, \dot{\epsilon}, p, T) = [\tau_a + \tau_i(\dot{\epsilon}, T) + \tau_e(\sigma_e, \dot{\epsilon}, T)] \frac{\mu(p, T)}{\mu_0}, \quad (1)$$

where σ_e is an evolving internal variable that has units of stress (also called the mechanical threshold stress), $\dot{\epsilon}$ is the strain rate, p is the pressure, T is the temperature, τ_a is the athermal component of the flow stress, τ_i is the intrinsic component of the flow stress due to barriers to thermally activated dislocation motion, τ_e is the component of the flow stress due to structure evolution (that is, strain hardening), μ is the shear modulus, and μ_0 is a reference shear modulus at 0 K and ambient pressure.

2.1. Athermal component. The athermal component of the yield stress is a function of grain size, dislocation density, distribution of solute atoms, and other long range barriers to dislocation motion. This component may either be constant or may evolve with deformation. A simple model for this component can be written as [Zerilli and Armstrong 1987; Banerjee 2007]:

$$\tau_a = \tau_0 + \frac{k}{\sqrt{d}}, \quad (2)$$

where τ_0 is the component due to far field dislocation interactions and the second term represents contribution due to the Hall–Petch effect where k is a material constant and d is the grain size.

2.2. Intrinsic component. In the MTS model, the intrinsic component of the flow stress is assumed to be dependent only on thermally activated mechanisms in the absence of solute diffusion. This component can be expressed in the phenomenological Arrhenius form given below (see [Kocks et al. 1975, Section 44], for an explanation)

$$\tau_i = S_i \sigma_i, \quad S_i := \left[1 - \left(\frac{k_b T}{g_{0i} b^3 \mu(p, T)} \ln \frac{\dot{\epsilon}_{0i}}{\dot{\epsilon}} \right)^{1/q_i} \right]^{1/p_i}, \quad (3)$$

where σ_i is the intrinsic component of the mechanical threshold stress (flow stress at 0 K), k_b is the Boltzmann constant, b is the magnitude of Burgers vector, g_{0i} is a normalized activation energy, $\dot{\epsilon}_{0i}$ is a constant reference strain rate, and p_i, q_i are constants. The constant p_i can take values between 0 and 1, while q_i can take values between 1 and 2.

In order to fit the high temperature data for 6061-T6 aluminum better, we have found that a better relation for τ_i has the form

$$\tau_i = \sigma_i \left[1 - \left\{ \left(\frac{k_b T}{g_{0i} b^3 \mu(p, T)} \ln \frac{\dot{\epsilon}_{0i}}{\dot{\epsilon}} \right)^{1/q_i} \right\}^2 \right]^{1/p_i}. \quad (4)$$

This relation is equivalent to using values of q_i between 0.5 and 1. If we use such a value in the original MTS model, the value of τ_i tends to become constant as the strain rate is increased. Recall that the original MTS model does not account for the effect of viscous drag at high strain rates, and the model predicts only a small increase in the flow stress with strain rate if q_i is between 1 and 2. The small reduction in the flow stress at high strain rates due to a value of q_i between 0.5 and 1 is easily offset by incorporating a simple model for viscous drag into the MTS model.

2.3. Viscous drag component. To incorporate viscous drag effects due to phonon damping, forest dislocations, and solute atoms, we add a term to the equation for τ_i of the form [Kumar and Kumble 1969;

Frost and Ashby 1971]

$$\tau_v = \frac{2}{\sqrt{3}} \frac{B}{\rho_m b^2} \dot{\epsilon}, \quad (5)$$

where B is the drag coefficient, ρ_m is the mobile dislocation density, and b is the magnitude of the Burgers vector. We assume that a good estimate of the value of the drag coefficient can be obtained if we consider only damping due to phonon scattering [Wolfer 1999]. However, at low temperatures (< 50 K), electron damping can dominate and a model for that should be used in addition to phonon damping (for more details see [Nadgorny 1988] p. 251–260).

The viscous drag coefficient can be written as [Kanel et al. 2001]

$$B \approx \lambda_p B_p = \frac{\lambda_p k_b T \omega_D^2}{\pi^2 \bar{c}^3}, \quad (6)$$

where B_p is the phonon drag coefficient, $\lambda_p \geq 1$ is a correction factor, ω_D is the Debye frequency and \bar{c} is the average sound speed. The Debye frequency is given by

$$\omega_D = \frac{2\pi k_b \theta_D}{h} = 2\pi \bar{c} \left(\frac{3\rho}{4\pi M} \right)^{1/3} = \left(\frac{6\pi^2 \rho}{M} \right)^{1/3} \bar{c}, \quad (7)$$

where θ_D is the Debye temperature, h is Planck's constant, and M is the atomic mass. The average sound speed is defined as

$$\frac{1}{\bar{c}^3} = \frac{1}{3} \left(\frac{2}{c_s^3} + \frac{1}{c_l^3} \right), \quad c_s = \sqrt{\frac{\mu}{\rho}}, \quad c_l = \sqrt{\frac{2\mu(1-\nu)}{\rho(1-2\nu)}}, \quad (8)$$

where c_s is the transverse sound speed, c_l is the longitudinal sound speed, and ν is the Poisson's ratio. Assuming that the Poisson's ratio is $1/3$, we can get an approximation for \bar{c} in terms of the shear sound speed as

$$\bar{c} \approx \left(\frac{24}{17} \right)^{1/3} c_s. \quad (9)$$

Alternatively, we can use Liebfried's model [Kumar and Kumble 1969; Brailsford 1970; Brailsford 1972; Wolfer 1999] for scattering of phonons by dislocations

$$B \approx \lambda_p B_p = \frac{\lambda_p q}{10c_s} \langle E \rangle, \quad (10)$$

where λ_p is a correction factor, q is the cross-section of the dislocation core per unit length and $\langle E \rangle$ is the average internal energy density. From Debye theory, we have

$$\langle E \rangle = \frac{3k_b T \rho}{M} D_3 \left(\frac{\theta_D}{T} \right), \quad \theta_D = \frac{h \bar{c}}{k_b} \left(\frac{3\rho}{4\pi M} \right)^{1/3}, \quad (11)$$

where the Debye function D_3 is defined as

$$D_3(x) := \frac{3}{x^3} \int_0^x \frac{z^3}{\exp(z) - 1} dz. \quad (12)$$

Note that the component of the flow stress due to viscous drag is both temperature and pressure dependent. A closed form expression can be obtained for the Debye integral in terms of polylogarithms (see for example [Truesdell 1945]). However, there is a singularity at $z = 0$ and analytic continuation methods are needed to evaluate the polylogarithms for values of $z > 0$. Alternatively, expansions in terms of Chebyshev polynomials [Ng et al. 1969; MacLeod 1996] may be used. For computational purposes, a look-up table of values (for example [Stegun 1972]) is the most efficient.

The second variable that is needed to compute τ_v is the density of mobile dislocations (ρ_m). The density of mobile dislocations depends on the prior strain history of a material and also on the strain, the temperature, and the strain rate. A simple model for the evolution of mobile dislocations as a function of plastic strain can be written as [Estrin and Kubin 1986; Barlat et al. 2002]

$$\begin{aligned} \frac{d\rho_m}{d\varepsilon_p} &= \frac{M_1}{b^2} \left(\frac{\rho_f}{\rho_m} \right) - I_2(\dot{\varepsilon}, T)\rho_m - \frac{I_3}{b}\sqrt{\rho_f}, \\ \frac{d\rho_f}{d\varepsilon_p} &= I_2(\dot{\varepsilon}, T)\rho_m + \frac{I_3}{b}\sqrt{\rho_f} - A_4(\dot{\varepsilon}, T)\rho_f, \end{aligned} \quad (13)$$

where ρ_f is the density of forest dislocations, b is the magnitude of the Burgers' vector, M_1 is a rate of mobilization of dislocations, I_2 is a rate of immobilization of mobile dislocations due to interactions between mobile dislocations, I_3 is another immobilization rate due to interactions with forest dislocations, and A_4 is the rate of annihilation of forest dislocations due to climb, cross-slip, or diffusion.

To solve the system of differential equations, Equation (13), we need initial values of ρ_m and ρ_f . We may also assume that the density of forest dislocations saturates at a value of ρ_{fs} , that is, $d\rho_f/d\varepsilon_p = 0$ at $\rho_f = \rho_{fs}$. Since the density of forest dislocations is easier to measure than the rate of annihilation A_4 , we use ρ_{fs} to obtain the following expression for A_4 :

$$A_4(\dot{\varepsilon}, T) = I_2(\dot{\varepsilon}, T) \frac{\rho_m}{\rho_{fs}} + \frac{I_3}{b} \frac{1}{\sqrt{\rho_{fs}}}. \quad (14)$$

The rate I_2 and the saturation value ρ_{fs} are, in general, temperature and strain-rate dependent. We assume that the strain rate dependence of these quantities is negligible for aluminum alloys and only consider temperature dependence. Based on the experimental observation that the temperature dependence of the flow stress of aluminum alloys follows a sigmoidal curve [Rosenberg et al. 1986], we assume that I_2 and ρ_{fs} have the following temperature dependence:

$$I_2(T) = I_{20} \left(\frac{\alpha_I}{1 + \exp[-\beta_I(1 - 2\hat{T})]} \right), \quad \rho_{fs}(T) = \rho_{fs0} \left(1 - \frac{\alpha_f}{1 + \exp[\beta_f(1 - 2\hat{T})]} \right), \quad (15)$$

where I_{20} is the value of I_2 at 0 K, ρ_{fs0} is the value of ρ_{fs} at 0 K, $\hat{T} = T/T_m$ where T_m is the melting temperature, α_I, α_f are scaling factors, and β_I, β_f are constants that have values greater than 3.

2.4. Structure evolution component. The evolution of dislocation density could be used to determine the component of the flow stress that depends on the evolution of the structure of a material (see for instance [Barlat et al. 2002]). However, Equations (13) form a stiff system of differential equations and are not ideal for numerical simulations, particularly explicit calculations needed for high rate processes.

Instead, we use the approach originally suggested by [Follansbee and Kocks \[1988\]](#) and describe the component of the flow stress due to structure evolution by

$$\tau_e(\sigma_e, \dot{\varepsilon}, T) = S_e \sigma_e, \quad S_e := \left[1 - \left(\frac{k_b T}{g_{0e} b^3 \mu(p, T)} \ln \frac{\dot{\varepsilon}_{0e}}{\dot{\varepsilon}} \right)^{1/q_e} \right]^{1/p_e}, \quad (16)$$

where σ_e is an internal variable (the structure evolution component of the mechanical threshold stress) that evolves with the deformation, g_{0e} is a normalized activation energy, $\dot{\varepsilon}_{0e}$ is a reference strain rate, and p_e, q_e are constants.

The evolution of σ_e is assumed to be governed by the modified Voce rule

$$\frac{d\sigma_e}{d\varepsilon_p} = \theta_0 \left[1 - \left(\frac{\sigma_e}{\sigma_{es}} \right)^2 \right], \quad (17)$$

where θ_0 is the initial hardening modulus and σ_{es} is the saturation value of σ_e . This equation can be solved in closed form along a constant temperature and strain-rate path to get

$$\sigma_e(\varepsilon_p) = \sigma_{es} \left[\frac{A_\sigma \exp(\beta) + 1}{A_\sigma \exp(\beta) - 1} \right] \quad A_\sigma := \frac{\sigma_{ei} + \sigma_{es}}{\sigma_{ei} - \sigma_{es}}, \quad \beta := \frac{2\theta_0 \varepsilon_p}{\sigma_{es}}, \quad (18)$$

where σ_{ei} is the value of σ_e at zero plastic strain and can be nonzero depending on the prior strain history of the material. For the special case where σ_{ei} is zero, we have

$$\sigma_e(\varepsilon_p) = \sigma_{es} \left[\frac{\exp(\beta) - 1}{\exp(\beta) + 1} \right]. \quad (19)$$

The initial hardening modulus is assumed to be given by a relation of the form

$$\theta_0 = \mu(p, T) \left[a_0 + a_1 \ln \left(\frac{\dot{\varepsilon}_\theta}{\dot{\varepsilon}} \right) + a_2 \widehat{T} + a_3 \widehat{T} \ln \left(\frac{\dot{\varepsilon}_\theta}{\dot{\varepsilon}} \right) \right], \quad (20)$$

where a_0, a_1, a_2, a_3 are constants and $\dot{\varepsilon}_\theta$ is a reference strain rate, and $\widehat{T} = T/T_m$ where T_m is the melting temperature. The value of θ_0 is not allowed to be negative.

The saturation value of structure evolution stress is computed using an Arrhenius type relation of the form

$$\ln \left(\frac{\sigma_{es}}{\sigma_{0es}} \right) = \left(\frac{k_b T}{g_{0es} b^3 \mu(\rho, T)} \right) \ln \left(\frac{\dot{\varepsilon}}{\dot{\varepsilon}_{0es}} \right), \quad (21)$$

where σ_{0es} is the saturation threshold stress for deformation at 0 K, g_{0es} is a normalized activation energy, and $\dot{\varepsilon}_{0es}$ is a reference strain rate.

2.5. Overdriven shock regime. At strain-rates between 10^9 s^{-1} and 10^{12} s^{-1} , the plastic wave overruns the elastic precursor and produces a steeper shock front than is attainable due to adiabatic elastic compression. Such a shock is called *overdriven*. In order to allow for overdriven shocks in our model, we

follow the approach used in the Preston–Tonks–Wallace model [Preston et al. 2003]. The flow stress under overdriven shock conditions (and assuming J_2 plasticity) is given by

$$\sigma_{ys} = \sigma_{ys0} + y_1 \left(\frac{\dot{\epsilon}}{\gamma \dot{\xi}} \right)^{y_2} \mu(p, T), \quad (22)$$

where σ_{ys0} is a lower limit on the flow stress, y_1 , y_2 are material parameters, γ is a scaling factor, and $\dot{\xi}$ is a reference strain rate. The reference strain rate is defined as

$$\dot{\xi} := \frac{1}{2} \left(\frac{4\pi\rho}{3M} \right)^{1/3} c_s, \quad (23)$$

where M is the atomic mass and c_s is transverse sound speed. We assume that there is no hardening in the overdriven shock regime.

2.6. Behavior after melting. After the temperature crosses the melting temperature (T_m), we have to model the material as a liquid. Though, strictly speaking, the behavior of an aluminum alloy melt is nonNewtonian, we use a Newtonian model of the following form to determine the deviatoric stress in the material after it has melted:

$$\sigma_y = \mu_v \dot{\epsilon}, \quad (24)$$

where μ_v is the shear viscosity of the melt. The shear viscosity of aluminum alloys can be estimated by first computing the viscosities of the constituent elements using the Andrade equation [da Andrade 1934; 1952] (alternative approaches can be found in [Kirkwood et al. 1949] and [Dinsdale and Quedsted 2004]). The viscosity of the alloy can then be calculated using the Moelwyn–Hughes model for binary alloys [Moelwyn-Hughes 1961; Dinsdale and Quedsted 2004]. However, since we are not interested in the details of the motion of the melt, we simplify our model by using a constant value of $\mu_v = 2.0 \times 10^{-4} \text{Pa} \cdot \text{s}$ in our calculations.

2.7. The extended flow stress model. The final expression for the flow stress is

$$\sigma_y = \begin{cases} \min \left\{ \left[\tau_v + (\tau_a + \tau_i + \tau_e) \frac{\mu}{\mu_0} \right], \sigma_{ys} \right\}, & \text{for } T < T_m, \\ \mu_v \dot{\epsilon}, & \text{for } T \geq T_m. \end{cases} \quad (25)$$

2.8. Shear modulus, melting temperature, EOS, specific heat. To complete the model, we need an expression for the shear modulus as a function of temperature and pressure. We have used a modified form of the Nadal–LePoac model [Nadal and Le Poac 2003] in which the pressure dependence of the shear modulus is determined by the generalized Guinan–Steinberg formula [Burakovsky and Preston 2005; Guinan and Steinberg 1975]:

$$\mu(p, T) = \frac{1}{\mathcal{F}(\widehat{T}, \zeta)} \left[\left\{ \mu_0 + p \frac{\partial \mu}{\partial p} \left(\frac{a_1}{\eta^{1/3}} + \frac{a_2}{\eta^{2/3}} + \frac{a_3}{\eta} \right) \right\} (1 - \widehat{T}) + \frac{\rho}{CM} k_b T \right], \quad (26)$$

$$\eta := \frac{\rho}{\rho_0}, \quad C := \frac{(6\pi^2)^{2/3}}{3} f^2,$$

where \mathcal{F} is a scaling factor that allows for a smooth transition of the shear modulus to melt, $\widehat{T} := T/T_m$, T_m is the melting temperature, ζ is a material parameter, μ_0 is the shear modulus at 0 K and ambient pressure, $\partial\mu/\partial p$ is the slope of the shear modulus versus pressure curve at 0 K, a_1, a_2, a_3 are material constants, ρ_0 is the mass density in the initial state, M is the atomic mass, f is the Lindemann constant, and

$$\mathcal{F}(\widehat{T}, \zeta) := 1 + \exp \left[-\frac{1 + 1/\zeta}{1 + \zeta/(1 - \widehat{T})} \right] \quad \text{for } \widehat{T} \in [0, 1 + \zeta]. \quad (27)$$

For situations where $\rho/\rho_0 < 0.95$ (high hydrostatic tension), we set the shear modulus to the value obtained for $\rho/\rho_0 = 0.95$.

We use the Burakovsky–Greeff–Preston (BGP) model [Burakovsky et al. 2003] to determine the melting temperature. This model improves upon the widely used Steinberg–Cochran–Guinan model [Steinberg et al. 1980] which is also based on the Lindemann criterion. In the BGP model, the Grüneisen gamma is modeled as

$$\Gamma(\rho) = \frac{1}{2} + \frac{\Gamma_1}{\rho^{1/3}} + \frac{\Gamma_2}{\rho^q}, \quad (28)$$

where Γ_1, Γ_2, q are material constants and $q > 1$. This form of $\Gamma(\rho)$ leads to the following expression for the melting temperature as a function of density:

$$T_m(\rho) = T_{m0}\eta^{1/3} \exp \left\{ 6\Gamma_1 \left(\frac{1}{\rho_0^{1/3}} - \frac{1}{\rho^{1/3}} \right) + \frac{2\Gamma_2}{q} \left(\frac{1}{\rho_0^q} - \frac{1}{\rho^q} \right) \right\}, \quad \eta := \frac{\rho}{\rho_0}, \quad (29)$$

where T_{m0} is the melt temperature when $\rho = \rho_0$.

The pressure is calculated using the Holzapfel equation of state [Holzapfel 1991a; 1991b; Burakovsky and Preston 2005] which has the form

$$p(\rho) = 3B_0 [\eta^{5/3} - \eta^{4/3}] \exp \left\{ 1.5(B'_0 - 3) [1 - \eta^{-1/3}] \right\}, \quad \eta := \frac{\rho}{\rho_0}, \quad (30)$$

where B_0 is the bulk modulus at $p = 0$ and B'_0 is the slope of the pressure-density curve at $p = 0$.

We use an empirical temperature-dependent specific heat model and ignore the difference between the values of specific heat at constant pressure and at constant volume. The specific heat model is of the form [Bhawalkar 2006]

$$C_v = \begin{cases} A_1 + B_1 T + C_1 T^2, & \text{for } T < 273 \text{ K}, \\ A_2 + B_2 T, & \text{for } T \geq 273 \text{ K}, \end{cases} \quad (31)$$

where A_1, A_2, B_1, B_2, C_1 are constants.

3. Evolution of temperature due to plastic dissipation

To determine the increase in temperature due to plastic dissipation, we first assume that the deformation gradient (\mathbf{F}) can be split multiplicatively into elastic and plastic parts ($\mathbf{F} = \mathbf{F}_e \cdot \mathbf{F}_p$). We also assume that there is no volume change during the plastic part of the deformation.

The elastic strain measure (\mathbf{E}_e) (with respect to the intermediate configuration) and the plastic strain rate (\mathbf{d}_p) are defined as (see [Wright 2002] for details)

$$\mathbf{E}_e := \frac{1}{2}[\mathbf{F}_e^T \cdot \mathbf{F}_e - \mathbf{1}], \quad (32)$$

$$\mathbf{d}_p := \frac{1}{2}[\mathbf{F}_e \cdot \dot{\mathbf{F}}_p \cdot \mathbf{F}_p^{-1} \cdot \mathbf{F}_e^{-1} + \mathbf{F}_e^{-T} \cdot \mathbf{F}_p^{-T} \cdot \dot{\mathbf{F}}_p^T \cdot \mathbf{F}_e^T]. \quad (33)$$

The stress measure in the intermediate configuration (that is, work conjugate to \mathbf{E}_e) is given by the pull-back of the Cauchy stress by the elastic part of the deformation gradient:

$$\tilde{\mathbf{S}} = \frac{\rho_0}{\rho} \mathbf{F}_e^{-1} \cdot \boldsymbol{\sigma} \cdot \mathbf{F}_e^{-T}. \quad (34)$$

Then the balance of energy can be expressed as

$$\rho T \dot{\eta} = -\nabla \cdot \mathbf{q} + \rho r + \boldsymbol{\sigma} : \mathbf{d}_p - \frac{\rho}{\rho_0} \sum_{n=1}^N Q_n \dot{q}_n, \quad (35)$$

where η is the entropy density, $\nabla \cdot (\bullet)$ denotes the divergence (with respect to the current configuration), r is a internal heat source term, $\boldsymbol{\sigma}$ is the Cauchy stress, and q_n is a set of internal variables. The quantities Q_n are work conjugate to the internal variables q_n and are defined as

$$Q_n := -\rho_0 \frac{\partial g}{\partial q_n}, \quad \text{for } n = 1 \dots N, \quad (36)$$

where $g(\tilde{\mathbf{S}}, T, q_n)$ is the specific Gibbs free energy (per unit mass) and ρ_0 is the density in the intermediate configuration (here assumed to be the same as that in the reference configuration).

The internal variable in the MTS model is σ_e , that is

$$q_1 = \sigma_e, \quad Q_1 = -\rho_0 \frac{\partial g}{\partial \sigma_e}. \quad (37)$$

Under adiabatic conditions which occur for high strain-rate deformations, the heat flux and internal heat sources can be neglected. In that case, the energy equation takes the form

$$\rho T \dot{\eta} = \boldsymbol{\sigma} : \mathbf{d}_p + \rho \frac{\partial g}{\partial \sigma_e} \dot{\sigma}_e. \quad (38)$$

Assuming that the entropy can be derived from the specific Gibbs free energy, it can be shown, after some algebra, that [Wright 2002]

$$\dot{\eta} = \frac{\partial \eta}{\partial \tilde{\mathbf{S}}} : \dot{\tilde{\mathbf{S}}} + \frac{\partial \eta}{\partial T} \dot{T} + \frac{\partial \eta}{\partial \sigma_e} \dot{\sigma}_e = \frac{1}{\rho_0} \frac{\partial \mathbf{E}_e}{\partial T} : \dot{\tilde{\mathbf{S}}} + \frac{C_v}{T} \dot{T} - \frac{1}{\rho_0} \frac{\partial \tilde{\mathbf{S}}}{\partial T} : \dot{\mathbf{E}}_e + \frac{\partial}{\partial T} \left(\frac{\partial g}{\partial \sigma_e} \right) \dot{\sigma}_e. \quad (39)$$

Substitution into the energy equation yields

$$\rho C_v \dot{T} = \boldsymbol{\sigma} : \mathbf{d}_p + \rho \left[\frac{\partial g}{\partial \sigma_e} - T \frac{\partial}{\partial T} \left(\frac{\partial g}{\partial \sigma_e} \right) \right] \dot{\sigma}_e. \quad (40)$$

It is conventional to use a factor χ (called the Taylor–Quinney coefficient) to relate the rate of change of temperature to the plastic work $\mathcal{P}_p := \boldsymbol{\sigma} : \mathbf{d}_p$ using the relation

$$\dot{T} = \frac{\chi \boldsymbol{\sigma} : \mathbf{d}_p}{\rho C_v}. \quad (41)$$

Therefore, from Equation (40), we observe that

$$\chi = 1 - \left[T \frac{\partial}{\partial T} \left(\frac{\partial g}{\partial \sigma_e} \right) - \frac{\partial g}{\partial \sigma_e} \right] \left(\frac{\rho \dot{\sigma}_e}{\boldsymbol{\sigma} : \mathbf{d}_p} \right). \quad (42)$$

Clearly χ evolves with σ_e and T , and cannot be assumed to be around 0.9 as is usually the case. However, the functional form of the Gibbs free energy function is not obvious and we have to make certain assumptions to obtain a function that matches experimental observations (see for instance [Hodowany et al. 2000]).

If we make the assumption that $\boldsymbol{\sigma} : \mathbf{d}_p \approx \sigma_y \dot{\varepsilon}_p$, the expression in Equation (42) reduces to

$$\chi = 1 - \left[T \frac{\partial}{\partial T} \left(\frac{\partial g}{\partial \sigma_e} \right) - \frac{\partial g}{\partial \sigma_e} \right] \left(\frac{\rho}{\sigma_y} \right) \frac{d\sigma_e}{d\varepsilon_p}.$$

Replacing the derivative of σ_e with Equation (17), we get

$$\chi = 1 - \left[T \frac{\partial}{\partial T} \left(\frac{\partial g}{\partial \sigma_e} \right) - \frac{\partial g}{\partial \sigma_e} \right] \left(\frac{\rho \theta_0}{\sigma_y} \right) \left[1 - \left(\frac{\sigma_e}{\sigma_{es}} \right)^2 \right]. \quad (43)$$

To model the specific Gibbs free energy $g(\tilde{\mathcal{S}}, T, \sigma_e)$, we follow the approach of Schreyer and Maudlin [2005] in postulating the following form for the expression involving the derivatives of g (which can be interpreted as a measure of the stored energy of cold work):

$$T \frac{\partial}{\partial T} \left(\frac{\partial g}{\partial \sigma_e} \right) - \frac{\partial g}{\partial \sigma_e} = F_1(T) F_2(\sigma_e), \quad (44)$$

where

$$F_1(T) = \frac{A_\chi}{1 + \exp[-B_\chi(1 - 2\hat{T})]}, \quad F_2(\sigma_e) = \frac{1}{\rho_0} \left(\beta_\chi + \frac{\sigma_e}{\sigma_{es}} \right)^{1/2}, \quad (45)$$

where $A_\chi > 1$, $B_\chi > 1$ are constants, $\hat{T} = T/T_m$, where T_m is the melting temperature, and $0 < \beta_\chi < 1$ is a factor that determines the value of χ at zero plastic strain. We can now calculate the rise in temperature due to plastic dissipation and use that to determine the amount of thermal softening due to plastic deformation. Note that a closed form expression for g that can be evaluated analytically cannot be derived from Equation (44) and further investigation into possible forms of g is needed.

4. Parameter estimation for 6061-T6 aluminum

Before the parameters of the extended MTS model for 6061-T6 aluminum can be estimated, we need the parameters of the shear modulus model. We also need the parameters of the associated melting

temperature model, the equation of state model, and the specific heat model. A detailed exploration of various models, their parameters, and comparisons with experimental data can be found in [Bhawalkar 2006]. The parameters of the models described by Equations (26), (27), (29), (30), and (31) are given in Appendix A, where we also show comparisons with experimental data of the predicted shear modulus, melting temperature, and specific heat.

4.1. Athermal component. In the ASM Handbook on nonferrous alloys [ASM 2002] we find that, for quasistatic tensile loading, at 644 K the yield strength of 6061-T6 aluminum is 12 MPa. The ratio μ/μ_0 for this temperature is 0.65. Hence, the maximum possible value of τ_a at 644 K is 18 MPa. Since this component is athermal, that value must be an upper bound on τ_a . It is possible that the upper bound on τ_a is even lower since the melting temperature of aluminum at ambient pressure is around 933 K. Therefore, it is reasonable to assume that $\tau_a \approx 10$ MPa for 6061-T6 aluminum.

4.2. Intrinsic component. To compute the intrinsic component of the flow stress (τ_i), we have to determine the parameters σ_{0i} and g_{0i} for a particular choice of $\dot{\epsilon}_{0i}$, p_i , and q_i (see Equation (4)). From (25) we see that in the low to moderate strain rate regime, the flow stress at zero plastic strain can be written as

$$\sigma_y = \tau_v + (\tau_a + \tau_i) \frac{\mu}{\mu_0}. \quad (46)$$

To determine τ_i , we need the viscous drag component τ_v and the experimentally determined initial yield stress σ_y . Recent studies [Ogawa 2001; Sakino 2006] show that for strain rates less than 10^4 s^{-1} , the viscous drag component τ_v for 6061-T6 aluminum is small compared to the thermally activated component τ_i . Because the experimental stress-strain data to which have access are for strain rates ranging from 10^{-5} s^{-1} to $8.0 \times 10^3 \text{ s}^{-1}$, we neglect τ_v during the process of computation of σ_i and g_{0i} .

Then, from Equations (4) and (46), we have

$$\left(\frac{\sigma_y - \tau_a}{\mu} - \frac{\tau_a}{\mu_0} \right)^{p_i} = \left(\frac{\sigma_i}{\mu_0} \right)^{p_i} - \left(\frac{\sigma_i}{\mu_0} \right)^{p_i} \left(\frac{1}{g_{0i}} \right)^{2/q_i} \left(\frac{k_b T}{b^3 \mu} \ln \frac{\dot{\epsilon}_{0i}}{\dot{\epsilon}} \right)^{2/q_i},$$

which is of the form $y = A - Bx^2$. A Fisher plot of the experimental data and the model fit is shown in Figure 1. Details of the procedure used to obtain the experimental data and the sources of the data can be found in [Bhawalkar 2006]. The best fit to the experimental data is obtained for $p_i = 0.75$, $q_i = 1.0$, and $\dot{\epsilon}_{0i} = 5.0 \times 10^8 \text{ s}^{-1}$. The Boltzmann constant was taken to be $k_b = 1.38 \times 10^{-23} \text{ J/K}$ and the magnitude of the Burgers' vector was assumed to be $b = 2.86 \times 10^{-10} \text{ m}$ [Wang et al. 1998].

The fit shown in the figure has an R^2 value of 0.6, an F -statistic of 60.7 and Student's t -values of 6.0 and 35.3 for the two coefficients. There are 81 data points. Using a t -value of 1.663 (corresponding to the 95% interval), we get the following range of values for σ_i and g_{0i} :

$$\begin{aligned} \sigma_i \text{ (low)} &= 343.7 \text{ MPa}, & \sigma_i \text{ (fit)} &= 366.6 \text{ MPa}, & \sigma_i \text{ (high)} &= 389.8 \text{ MPa}, \\ g_{0i} \text{ (low)} &= 0.51, & g_{0i} \text{ (fit)} &= 0.59, & g_{0i} \text{ (high)} &= 0.71. \end{aligned}$$

4.3. Viscous drag component. The viscous drag component of the flow stress is given by Equation (5). Two quantities have to be determined before τ_v can be computed: the drag coefficient B and the mobile dislocation density ρ_m .

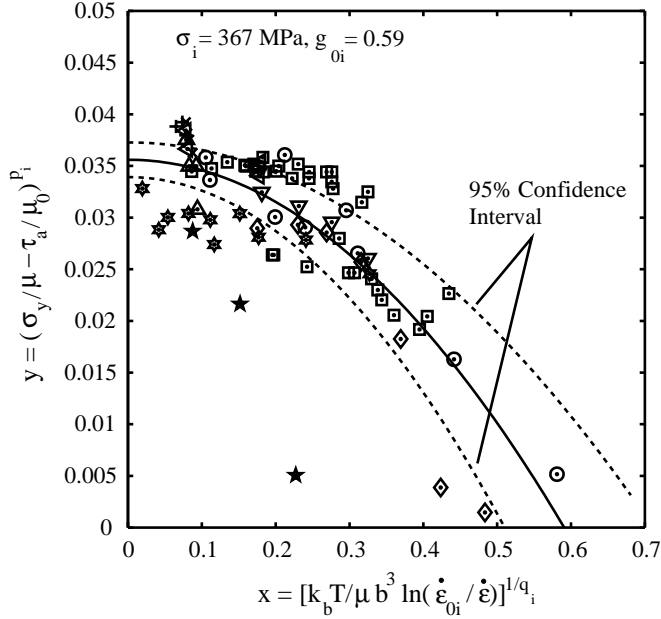


Figure 1. Fisher plot used to determine σ_i and g_{0i} for 6061-T6 aluminum alloy. The solid line shows the model fit to the data. The dotted lines show the range of the model for a confidence level of 95. The symbols represent experimental data. The sources of the experimental data are \circ [Hoge 1966]; \square [Green and Babcock 1966; Eleiche 1972]; ∇ [Davidson 1973]; \star [Rosenberg et al. 1986]; \triangle [Yadav et al. 1995]; \diamond [Brown et al. 1996]; \times [Lee et al. 2000]; $*$ [Ogawa 2001]; $<$ [Cady and Gray III 2001; Fish et al. 2005]; $+$ [Lesuer et al. 2001; Jia and Ramesh 2004].

The drag coefficient can be estimated using either (6) or (10). Figure 2 shows the phonon drag coefficients predicted by the two models. The constants used for the models are:

$$\begin{aligned}
 k_b &= 1.38 \times 10^{-23} \text{ J/K}, & b &= 2.86 \times 10^{-10} \text{ m}, \\
 M &= 26.98 \text{ amu}, & \rho_0 &= 2700 \text{ kg/m}^3, \\
 h &= 6.62 \times 10^{-34} \text{ kg} \cdot \text{m}^2/\text{s}, & q &= 3b.
 \end{aligned}$$

Values of the Debye function (D_3) were calculated using an algorithm given in [MacLeod 1996]. As the pressure increases there is a decrease in B_p . However, the change is small and does not significantly affect the value of τ_v . At temperatures below 50 K, electron drag effects become dominant. Note that this is not reflected by the model in Equation (10). However, since that model predicts a lower value of B_p at high temperatures, we have used (10) in subsequent calculations.

The evolution of dislocation density is harder to quantify. We have chosen parameters for Equation (13) in such a way that, at high strain rates, the viscous drag component of the flow stress approximates

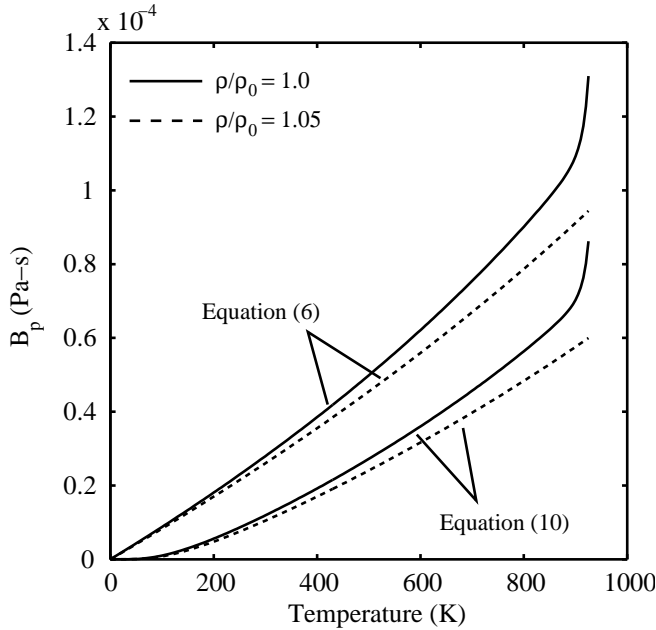


Figure 2. Phonon drag coefficient (B_p) as a function of temperature. The upper curve shows the prediction of the model in Equation (6) and the lower curve shows the values predicted by the model in (10).

experimentally determined yield stress at room temperature. The parameters used for the model are:

$$\begin{aligned}
 b &= 2.86 \text{ m}^{-2}, & M_1 &= 4.14 \times 10^{-7}, & I_{20} &= 5.53 \times 10^3, \\
 I_3 &= 3.81 \times 10^{-2}, & \alpha_I &= 1.0, & \beta_I &= 8.0, \\
 \rho_{f,s0} &= 1.0 \times 10^{14} \text{ m}^{-2}, & \alpha_f &= 0.4, & \beta_f &= 4.0.
 \end{aligned}$$

Using initial values of $\rho_m = 1.0 \times 10^9 \text{ m}^{-2}$ and $\rho_f = 1.0 \times 10^{10} \text{ m}^{-2}$, we get the evolution of the dislocation densities shown in Figure 3. A constant density of $2700 \text{ kg}\cdot\text{m}^{-3}$ has been used in the computation.

The saturation value of ρ_m is approximately $2 \times 10^{11} \text{ m}^{-2}$ at room temperature, which is three orders of magnitude lower than that suggested by Estrin and Kubin [1986]. However, higher values of ρ_m lead to unreasonably low values of the viscous drag component of the flow stress. Also, similar low values of ρ_m are suggested by Kumar and Kumble [1969]. Note that the value of ρ_f is close to the accepted value of total dislocation density in aluminum and its alloys, on the order of $1.0 \times 10^{14} \text{ m}^{-2}$. It can also be observed that the density of forest dislocations decreases with temperature while that of mobile dislocations increases with temperature. Also, our model indicates that the rate of annihilation of forest dislocations is greater than the rate of generation of mobile dislocations as the temperature increases (see Figure 3).

The viscous drag component (τ_v) of the flow stress can now be computed using (5). Figure 4 shows the variation of τ_v with strain rate and temperature (for a plastic strain of 0.2). We have used a value of $\lambda_p = 1$ in the calculations. From the figure it is clear that the viscous drag becomes significant only beyond strain rates of 10^5 s^{-1} . Also, drag effects appear to be negligible at low temperatures.

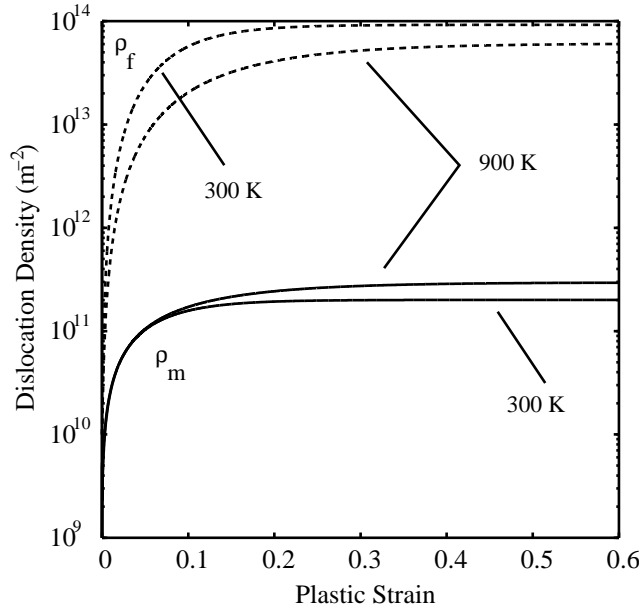


Figure 3. Evolution of dislocation density as a function of plastic strain and temperature. The solid curves show the evolution of ρ_m while the dotted curves show the evolution of ρ_f .

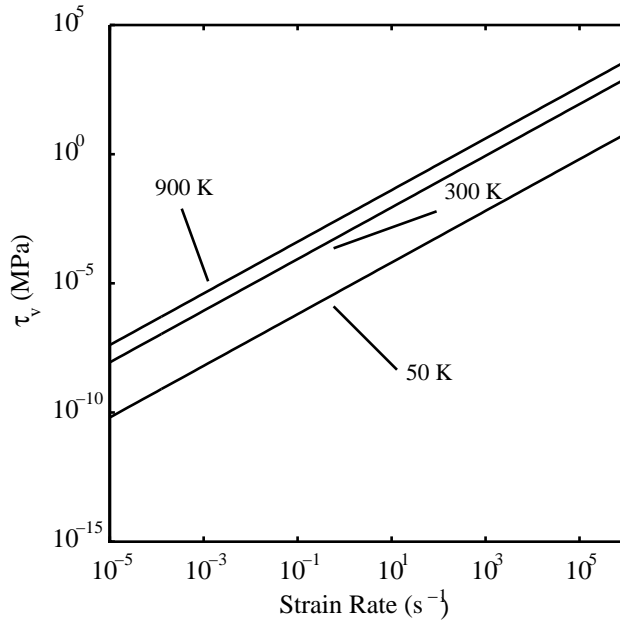


Figure 4. Variation of the viscous drag component of the flow stress of 6061-T6 aluminum with strain rate and temperature for a plastic strain of 0.2.

4.4. Structure evolution component. To determine the structure evolution component (τ_e) of the flow stress we need to compute the evolution of the internal variable σ_e using equation Equation (19). This equation requires two variables: the saturation value σ_{es} of σ_e and the initial hardening rate θ_0 .

To compute σ_{es} using (21), we need the parameters σ_{0es} , g_{0es} , and $\dot{\epsilon}_{0es}$. Given a set of true stress versus true plastic strain curves, the first step in the process of determination of these parameters is to compute the temperature at each point on the curve (for experimental data that are not given in isothermal form). The temperature is computed using (41) with the value of χ being obtained using:

$$\chi = \min\{[0.019 - 0.089 \ln(\varepsilon_p) + 9.23\varepsilon_p^2], 0.9\}, \quad (47)$$

where ε_p is the true plastic strain. Equation (47) is a least squares fit to the experimental data for 2024-T351 aluminum given in [Ravichandran et al. 2001]. The experimental σ_e versus true plastic strain curves are then computed using the relation

$$\sigma_e = \frac{1}{S_e} \left(\sigma_y \frac{\mu_0}{\mu} - \tau_a - \tau_i \right).$$

Note that the quantity τ_i for each curve is the value used to compute the Fisher plot in Figure 1, unlike the procedure used elsewhere [Goto et al. 2000; Banerjee 2007]. The parameters used to compute S_e (see (16)) are:

$$\begin{aligned} k_b &= 1.38 \times 10^{-23} \text{ J} \cdot \text{K}^{-1}, & b &= 2.86 \times 10^{-10} \text{ m}^{-2}, & \dot{\epsilon}_{0e} &= 10^9 \text{ s}^{-1}, \\ g_{0e} &= 3.0, & p_e &= 0.75, & q_e &= 1.0. \end{aligned}$$

The density is assumed to have a constant value of $2700 \text{ kg} \cdot \text{m}^{-3}$ during deformation. Plots of σ_e versus ε_p for all the stress-strain curves used in this work can be found in [Bhawalkar 2006].

After the σ_e versus ε_p curves have been computed, a quadratic polynomial is fit to each curve. The saturation value of σ_e and the temperature at saturation are computed by determining the strain at which the slope of the quadratic curve becomes zero. A Fisher plot can now be drawn to determine the values of σ_{0es} and g_{0es} by writing Equation (21) in the form

$$\ln(\sigma_{es}) = \ln(\sigma_{0es}) - \frac{k_b T}{g_{0es} b^3 \mu} \ln \left(\frac{\dot{\epsilon}_{0es}}{\dot{\epsilon}} \right).$$

To compute the Fisher plot we assume that $\dot{\epsilon}_{0es} = 10^8 \text{ s}^{-1}$ and get the fit shown in Figure 5. The statistics of the fit are: $R^2 = 0.41$, F -statistic = 36.2, t -statistics = 6.0 and 157. There are 52 sample points. Therefore, though the fit is poor, it is statistically significant. The range of values of σ_{0es} and g_{0es} from the fit that are within the 95% confidence interval (corresponding to a t value of 1.675) are:

$$\begin{aligned} \sigma_{0es} \text{ (low)} &= 122.3 \text{ MPa}, & \sigma_{0es} \text{ (fit)} &= 149.5 \text{ MPa}, & \sigma_{0es} \text{ (high)} &= 182.7 \text{ MPa}, \\ g_{0es} \text{ (low)} &= 0.235, & g_{0es} \text{ (fit)} &= 0.30, & g_{0es} \text{ (high)} &= 0.416. \end{aligned}$$

To compute the initial hardening rate (θ_0), we first compute the slopes (θ) of the σ_e versus ε_p curves at each point on the curve. The slope θ is then plotted against $1 - (\sigma_e/\sigma_{es})^2$ and a straight line with zero intercept is fitted to the resulting curve. This model gives a better fit to the experimental data than the more commonly used hyperbolic tangent hardening rule [Follansbee and Kocks 1988; Goto et al. 2000;

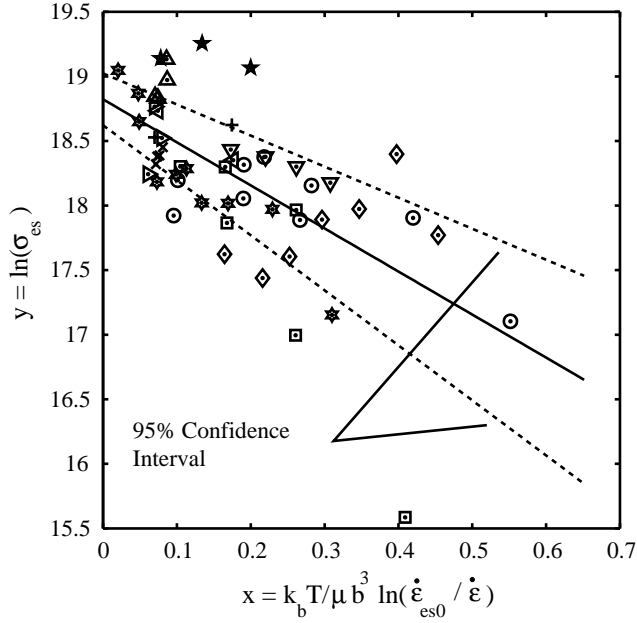


Figure 5. Fisher plot used to determine σ_{0es} and g_{0es} for 6061-T6 aluminum alloy. The solid line shows the model fit to the data. The dotted lines show the range of the model for a confidence level of 95%. The symbols represent experimental data. The sources of the experimental data are \circ [Hoge 1966]; \square [Green and Babcock 1966; Eleiche 1972]; ∇ [Davidson 1973]; \star [Rosenberg et al. 1986]; \triangle [Yadav et al. 1995]; \diamond [Brown et al. 1996]; \times [Lee et al. 2000]; $*$ [Ogawa 2001]; $+$ [Cady and Gray III 2001]; $<$ [Fish et al. 2005]; $>$ [Lesuer et al. 2001; Jia and Ramesh 2004].

[Banerjee 2007]. The slope of the straight line fit gives us the value of θ_0 for each curve. A least squares surface fit of θ_0 as a function of strain rate and temperature (assuming that the parameter $\dot{\epsilon}_\theta = 10^{10} \text{ s}^{-1}$) gives us the following parameters for Equation (20):

$$\begin{aligned} a_0 &= 0.147, & a_1 &= -2.4 \times 10^{-3}, \\ a_2 &= -0.14, & a_3 &= 3.3 \times 10^{-3}. \end{aligned}$$

The spread of the data is quite large and the R^2 value for the fit is 0.38. The F -statistic for the fit is 3.2 which implies that the fit may not be statistically significant. However, comparisons with experimental data show that the predicted flow stresses are reasonable when we use the above parameters. For the details of an alternative approach for modeling the hardening of 6061-T6 aluminum in the context of the MTS model see [Bhawalkar 2006].

4.5. Overdriven shock regime. In the overdriven shock regime, we assume that the flow stress is given by Equation (22) and that the material does not harden. The parameters of the model are determined by fitting a curve to experimental data on 1100 aluminum and 2024 aluminum from [Tonks 1993]. We assume that 6061-T6 aluminum also shows a similar behavior. The fit to the data is shown in Figure 6.

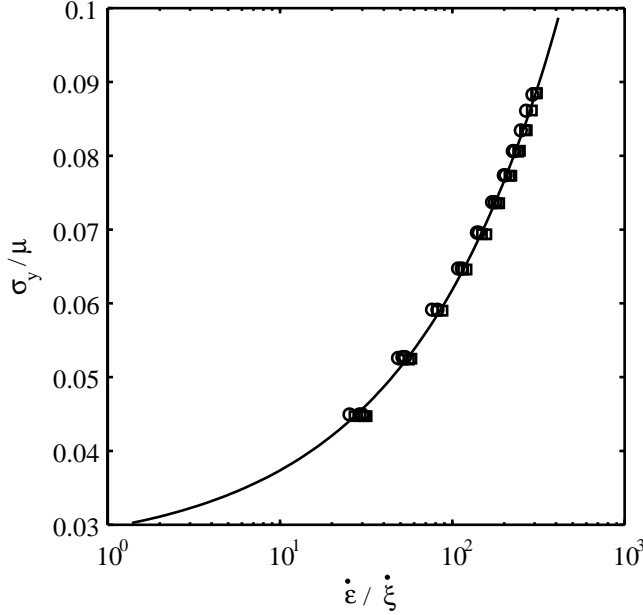


Figure 6. Comparison of the yield stress predicted by the overdriven shock model, Equation (22), with experimental data. The solid line shows the predicted values. The circles show experimental data for 1100 aluminum alloy while the squares show experimental data for 2024 aluminum alloy.

The following parameters have been used in the calculation:

$$\begin{aligned} \rho_0 &= 2700 \text{ kg} \cdot \text{m}^{-3}, & \rho &= 2710 \text{ kg} \cdot \text{m}^{-3}, & T &= 298 \text{ K}, \\ \sigma_{ys0} &= 700 \text{ MPa}, & M &= 26.98 \text{ amu}, & \gamma &= 0.0001. \end{aligned}$$

The parameters determined from the fit are: $y_1 = 3.54 \times 10^{-3}$ and $y_2 = 0.5$.

4.6. Taylor–Quinney coefficient. The Taylor–Quinney coefficient (χ) can be computed using Equation (43). To evaluate this equation we need the parameters A_χ , B_χ , and β_χ in (45). We have determined these parameters by comparing the predictions of our model with experimental data for 2024-T3 aluminum from [Hodowany et al. 2000]. Figures 7 (left) and 7 (right) show the predicted values of χ for initial temperatures of 300 K and 500 K, respectively. The parameters used for the plots are: $A_\chi = 0.2$, $B_\chi = 12$, and $\beta_\chi = 0.3$. A constant mass density of 2700 kg m^{-3} was used for the calculations. Note that these parameters will change if θ_0 or σ_{es} are changed.

From Figure 7 (left) we observe that, at a temperature of 300 K, the values of χ are nearly the same for strain rates between 0.001 s^{-1} to 3000 s^{-1} . This is similar to the experimental observations in [Hodowany et al. 2000]. However, the value of χ increases as the flow stress increases with increasing strain rate due to viscous drag effects. This effect is similar to that observed in rate-dependent α -titanium in [Hodowany et al. 2000]. Hence, our model behaves in a reasonable manner at room temperature. On the other hand, the experimental data for 2024-T3 aluminum indicates that low values of χ (less than 0.5) are obtained up to a plastic strain of around 0.3 while our model predicts such values only up to a plastic strain of

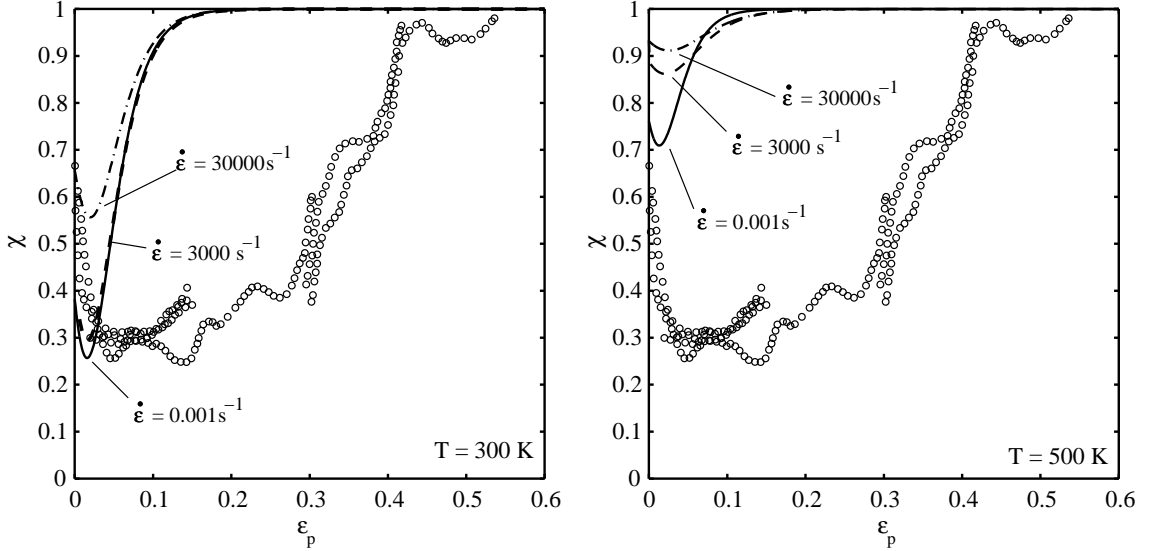


Figure 7. Variation of the Taylor–Quinney coefficient (χ) with plastic strain and strain rate: $T = 300$ K (left), $T = 500$ K (right). The circles show experimental data for 2024-T3 aluminum alloy from [Hodowany et al. 2000] for a strain rate of 3000 s^{-1} and at 300 K. The solid lines represent predicted values for a strain rate of 0.001 s^{-1} . The dashed lines represent predicted values for a strain rate of 3000 s^{-1} and the dash-dot lines represent predictions for a strain rate of $30,000 \text{ s}^{-1}$.

0.05. The width of this zone depends on the initial hardening modulus (θ_0) and on the saturation stress (σ_{es}). The discrepancy between our model for 6061-T6 aluminum and the experimental data for 2024-T3 aluminum indicates that θ_0 for 2024-T3 aluminum might be lower than that for 6064-T6 aluminum. It is also possible that the saturation stress computed by our model is lower than should be expected.

At 500 K (see Figure 7 (right)), the value of χ is considerably higher than at 300 K. We do not have any experimental data to verify that this behavior is reasonable. Experiments at high temperatures are needed to arrive at the correct form of the function $F_1(T)$ in Equation (44). As designed, our the model predicts values of χ close to 1.0 at high temperatures. However, experiments are needed to verify that this is indeed the case for metals and that the stored energy approaches zero at high temperatures.

5. Comparisons of model with experimental data

The verification of the extended MTS flow stress model presented in this paper can be pursued in a number of ways. The strain rate dependence of the model can be verified by computing the flow stress as a function of strain rate (at a constant temperature and plastic strain) and comparing the predicted values with experimental data. A similar comparison of the flow stress as a function of temperature (with the strain rate kept constant) can be used to verify the temperature dependence of the model. Direct comparisons with uniaxial stress-strain curves can also be performed for various strain rates and temperatures.

Numerical simulation is necessary for a more comprehensive validation of the model. Examples of such numerical validation tests have been provided in [Bhawalkar 2006]. Details of the numerical algorithm and various validation tests will be discussed in a forthcoming publication.

5.1. Flow stress as a function of strain rate. Figure 8 shows a comparison of the flow stress predicted by the extended MTS model with experimental data (as a function of strain rate). In the computations performed, the evolution equations for dislocation density were not solved because of the excessive computational cost involved. Rather, a constant value of $\rho_m = 2.0 \times 10^{11} \text{ m}^{-2}$ and a scaling factor of $\lambda_p = 1.0$ were used to compute the component of the flow stress due to viscous drag (τ_v). Adiabatic heating is assumed for strain rates greater than 100 s^{-1} .

Experimental data from [Yadav et al. 1995; Jia and Ramesh 2004; Sakino 2006] for 6061-T6 aluminum alloy suggest that viscous drag effects are small for strain rates up to $3 \times 10^4 \text{ s}^{-1}$. This is in contrast with data given in [Lesuer et al. 2001] which suggests that drag effects become important at strain rates of 10^4 s^{-1} or less. Sakino [2006] has shown quite convincingly that drag effects become important at lower strain rates only in the annealed alloy 6061-O aluminum but not in the hardened 6061-T6 aluminum alloy. Our model slightly overestimates the flow stress for strain rates less than 10^5 s^{-1} but matches quite

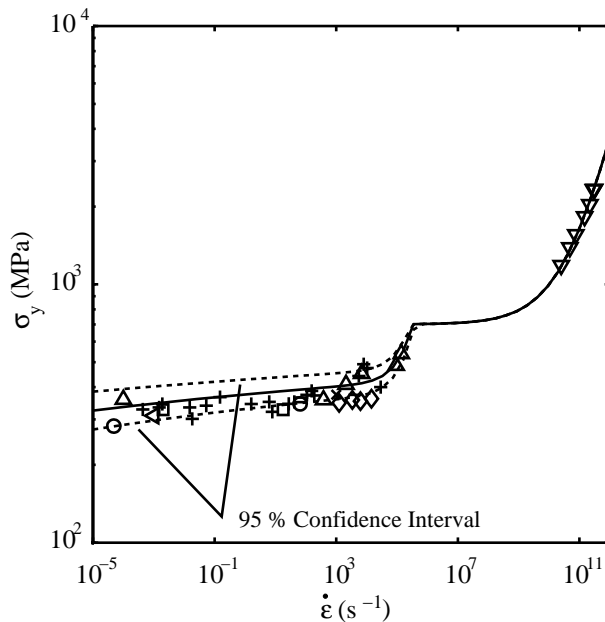


Figure 8. The flow stress of 6061-T6 aluminum alloy as a function of strain rate (at 298 K and a plastic strain of 0.2). The solid line shows the prediction of the extended MTS model. The dashed lines show the range of predicted flow stresses in the 95 % confidence interval. The symbols represent experimental data. The sources of the experimental data are \circ [Hoge 1966]; \square [Green and Babcock 1966; Eleiche 1972]; ∇ [Tonks 1993]; \triangle [Yadav et al. 1995]; \times [Lee et al. 2000]; \triangleleft [Cady and Gray III 2001]; $+$ [Lesuer et al. 2001; Jia and Ramesh 2004]; \diamond [Sakino 2006].

well with the available data for higher strain rates. The 95% confidence interval shown in the figure is based on the low and high values of σ_i , g_{0i} , σ_{0es} , and g_{0es} discussed earlier.

From Figure 8 we can also observe a change in slope of the flow stress-strain rate curve at a strain rate of around 10^6 s^{-1} . Experimental data are not available for the strain rate regime between 10^6 s^{-1} and 10^{10} s^{-1} . The curves for the overdriven shock regime and the moderate strain rate regime can be matched only if there is a change in slope as shown in the figure. A better model for the intermediate strain rate regime can only be determined after experiments (or possibly numerical simulations) have been performed for this regime. In the model that we have presented in this paper, we assume that the viscous drag component of the flow stress saturates when σ_y reaches a value of 700 MPa and any additional increase in the flow stress is due to shock effects as discussed in [Wallace 1981a; 1981b].

5.2. Flow stress as a function of temperature. Figure 9 shows comparisons between experimental data and the flow stress predicted by our model as a function of temperature. At low strain rates (see Figure 9, left), the decrease in flow stress with temperature is predicted quite accurately by the model. Near the melting temperature, the contributions of τ_i and τ_e are close to zero and the flow stress is determined by the athermal component τ_a . Beyond the melting temperature, the stress is computed by the Newtonian fluid model in Equation (24) and the stress is quite small at low strain rates. Adiabatic heat is assumed for the high strain rate calculations.

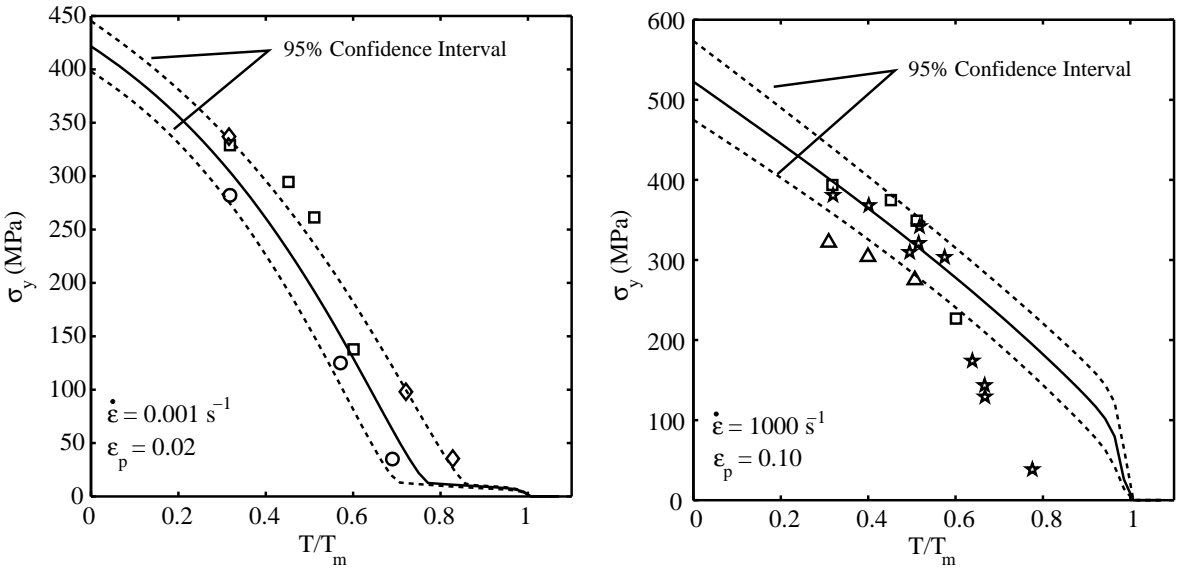


Figure 9. The flow stress of 6061-T6 aluminum alloy as a function of temperature. Left: strain rate = 0.001 s^{-1} and plastic strain = 0.02. Right: strain rate = 1000 s^{-1} and plastic strain = 0.1. The solid lines show the prediction of the extended MTS model. The dashed lines show the range of values predicted if we use the low or high values of the parameters σ_i , g_{0i} , σ_{0es} , and g_{0es} . The symbols represent experimental data. The sources of the experimental data are: \circ [Hoge 1966]; \square [Green and Babcock 1966; Eleiche 1972]; \diamond [Dike et al. 1997]; \star [Rosenberg et al. 1986]; \triangle [Ogawa 2001].

At a strain rate of 1000 s^{-1} , our model predicts the flow stress quite well up to $T/T_m = 0.6$ (see Figure 9, right). At higher temperatures, the flow stress is overestimated. The prediction of our model is reasonable if we consider the full set of data used to arrive at the parameters. It is expected that the predicted temperature dependence of our model can be improved at more experimental data at high temperatures and high strain rates become available.

5.3. Flow stress as a function of pressure. Figure 10 shows the predicted pressure dependence of the flow stress of 6061-T6 aluminum at a strain rate of 0.001 s^{-1} . The increase in flow stress with increasing pressure is underestimated by our model. Since the pressure dependence of the flow stress at low strain rates is determined by the pressure dependence of the shear modulus, the figure shows that the pressure dependence of the shear modulus is not enough to account for the pressure dependence of the flow stress. We also observe that

$$\frac{\partial \sigma_y}{\partial p} \approx 2 \frac{\partial \mu}{\partial p}.$$

This suggests that the pressure dependence of the flow stress in the Steinberg–Cochran–Guinan model [Steinberg et al. 1980] (which is used widely for high strain rate calculations and assumes that $\partial \sigma_y / \partial p \approx \partial \mu / \partial p$) is not correct. We have also observed a similar behavior for OFHC copper [Banerjee 2005a]. An explicit pressure dependence of the initial hardening modulus and the saturation stress is needed to

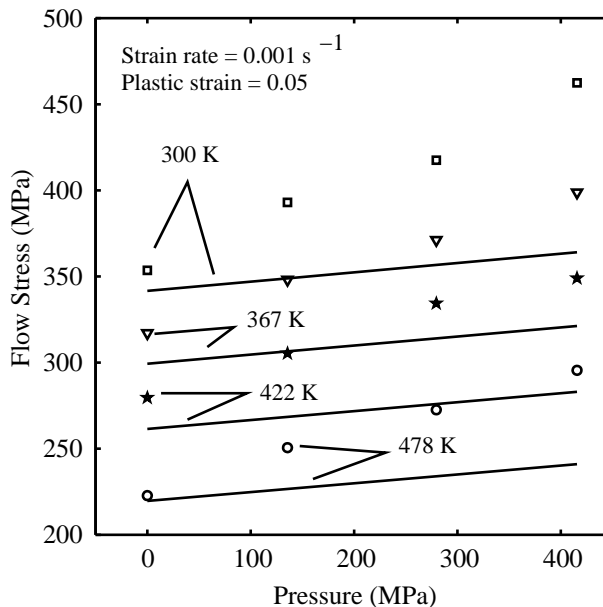


Figure 10. The flow stress of 6061-T6 aluminum alloy as a function of pressure for a strain rate of 0.001 s^{-1} and a plastic strain of 0.05. The solid lines show the predictions of the extended MTS model. The symbols represent experimental data from [Davidson 1973] at the following temperatures range of values predicted if we use the low or high values of □ 300 K, ▽ 367 K, ★ 422 K, and ○ 478 K.

model the plastic behavior of metals under hydrostatic pressure. Such a dependence has to be based on physical considerations and needs further research.

5.4. Uniaxial stress-strain curves. The predicted variation of the flow stress with plastic strain at a strain rate of 0.001 s^{-1} is shown in Figure 11. The calculations were performed using a constant mass density of $2700 \text{ kg}\cdot\text{m}^{-3}$, and a constant mobile dislocation density of $2 \times 10^{11} \text{ m}^{-2}$. Isothermal conditions were assumed.

Figure 11 (left) shows the predicted flow stress at ambient pressure. The flow stress predicted by our model tends to saturate at a lower value than seen in the experimental data. Also the predicted initial hardening modulus is lower than that suggested by the experiments at 367 K and 422 K and higher than experiment at 589 K. However, the initial yield stress predicted by our model is remarkably close to that seen in experiments considering the experimental variability that is observed (for further data on experimental variability at a given temperature and strain rate see [Bhawalkar 2006]). The experimental stress-strain curves show strain softening. A damage model that includes void nucleation and growth can be used to predict that portion of the stress-strain curve. For the sake of simplicity, we do not discuss damage models and their implications in this paper.

Figure 11 (right) shows the predicted flow stress at a pressure of 276 MPa. We notice that the initial yield stress is predicted quite accurately by our model at this pressure for all the temperatures shown in the figure. However, the hardening rate and the saturation stress are underestimated. The error decreases with

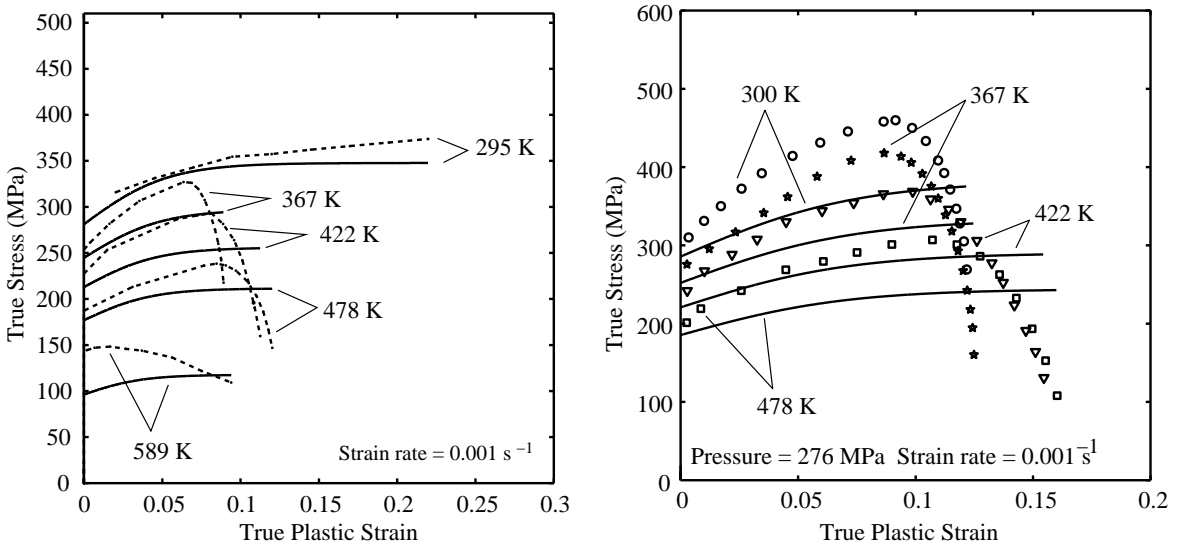


Figure 11. The flow stress of 6061-T6 aluminum alloy as a function of true plastic strain at a strain rate of 0.001 s^{-1} . Left: Ambient pressure; right: pressure = 276 MPa. The solid lines show the prediction of the extended MTS model. On the left side of the figure the dashed lines represent the experimental data. On the right, the symbols represent the experimental data. The data at 295 K are from [Fish et al. 2005]. The data at 300 K, 367 K, 422 K, and 478 K are from [Davidson 1973]. The data at 589 K are from [Green and Babcock 1966; Eliche 1972].

increasing temperature. Similar plots at higher pressures show that the error increases with increasing pressure at a particular temperature (see also Figure 10). As mentioned before, an explicit pressure dependence of the initial hardening rate and the saturation stress is needed to improve our model.

Figure 12 shows the predicted stress-strain curves for strain rates greater than 1000 s^{-1} . The change in mass density due to large pressures that develop at such strain rates has been neglected in our computations. To keep the computations efficient, we have not used the evolution equations for dislocation density and have instead used a constant value of $\rho_m = 2 \times 10^{11} \text{ m}^{-2}$. Adiabatic heating and the resulting thermal softening has been assumed for all these computations.

From Figure 12 (left) we observe that the flow stress predicted by the extended MTS model matches well with the experimental data for temperatures of 77 K and 298 K. At 473 K, the predicted flow stress is higher than experiment and does not capture the strain softening due to accumulating damage in the material. At 618 K, the predicted flow stress is considerably higher than experiments suggest. As observed before, our model does not predict enough thermal softening of 6061-T6 aluminum at higher temperatures— particularly at high strain rates. It is important to note here that 6061-T6 aluminum is a precipitation hardened alloy containing magnesium and silicon as alloying elements. At about 550 K, the solute molecules start to diffuse toward the grain boundaries and the alloy softens considerably. Though the extended MTS model is an improvement over the original MTS model (see [Bhawalkar

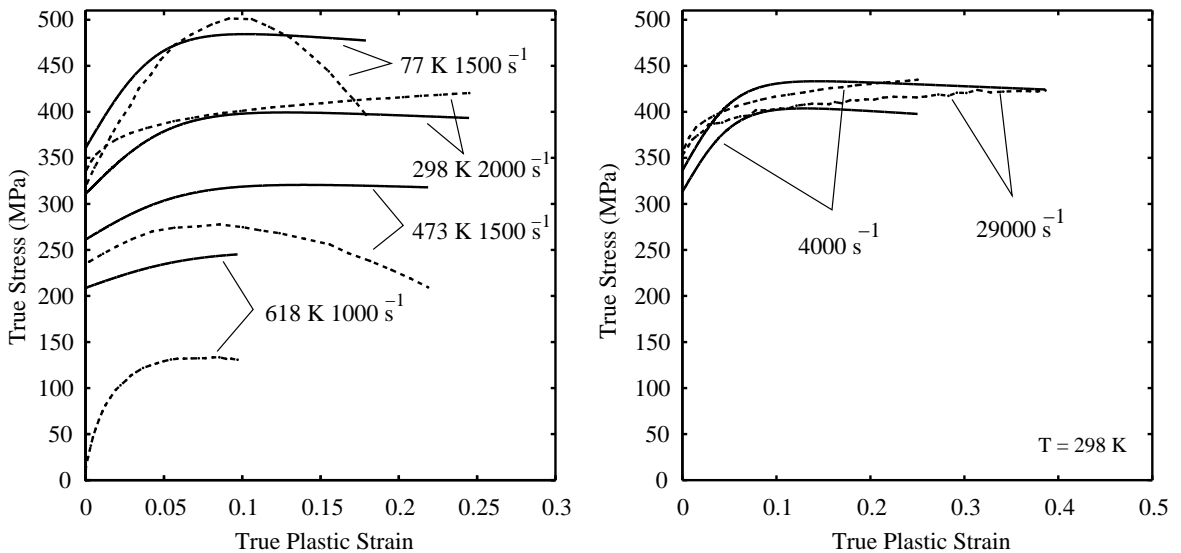


Figure 12. The flow stress of 6061-T6 aluminum alloy as a function of plastic strain at moderately high strain rates. Left: strain rate = $1000\text{--}2000 \text{ s}^{-1}$. Right: temperature = 298 K. The solid lines show the prediction of the extended MTS model. The dashed lines represent the experimental data. The sources of the experimental data are: strain rate = 1000 s^{-1} from [Rosenberg et al. 1986], strain rate = 1500 s^{-1} from [Ogawa 2001], strain rates = 2000 s^{-1} and 4000 s^{-1} from [Lee et al. 2000], strain rate = 29000 s^{-1} from [Jia and Ramesh 2004].

2006] for similar results with the original MTS model), the amount of thermal softening continues to be overestimated by the extended model.

Figure 12 (right) shows that the experimental data does not show any significant strain rate dependence at room temperature (at least up to a strain rate of $29,000 \text{ s}^{-1}$). Our model predicts a small increase in flow stress with strain rate and some thermal softening at strains greater than 0.2. However, the overall match with experiment is quite good. Also the initial rate of hardening appears to be lower than that predicted. However, stress-strain data from split Hopkinson pressure bar experiments are suspect for strains less than 0.1 and conclusions cannot be drawn about the accuracy of our hardening model from this figure.

6. Model predictions

In this section we discuss the predictions of our flow stress model for a large range of conditions. The mass density is assumed to be constant ($2700 \text{ kg}\cdot\text{m}^{-3}$) in our calculations. Since the phonon drag coefficient increases with temperature, assuming a constant mobile dislocation density for all temperatures gives unrealistic flow stresses at strain rates greater than 10^4 s^{-1} . On the other hand, using the evolution equations to determine ρ_m is not computationally efficient. Hence we have used the following equation to determine ρ_m for our calculations in this section:

$$\rho_m \approx \rho_{m0}(1 + \widehat{T})^m,$$

where ρ_{m0} is the dislocation density at 0 K, $\widehat{T} = T/T_m$ where T_m is the melting temperature, and m is a constant. The values of these parameters that we have used are: $\rho_{m0} = 1.3494 \times 10^{11} \text{ m}^{-2}$ and $m = 1.4114$. Using these parameters gives us a value of $\rho_m = 2 \times 10^{11} \text{ m}^{-2}$ at 300 K and $\rho_m = 3.5 \times 10^{11} \text{ m}^{-2}$ at 900 K. The value of ρ_m is not allowed to vary with plastic strain.

Figure 13 (left) shows the flow stress predicted by our model as a function of strain rate at a plastic strain of 0.2 and for a range of temperatures. From the figure we observe that at low strain rates, the flow stress is strongly dependent on temperature. As the strain rate reaches around 10^5 s^{-1} , the model predicts that the temperature dependence of the flow stress decreases until the temperature has hardly any effect.

Figure 13 (right) shows the flow stress as a function of strain rate at a temperature of 400 K and for various levels of strain. In this case we observe that the flow stress saturates rather rapidly (at a plastic strain of around 0.2). This indicates that the material exhibits little strain hardening. However, simulations of Taylor impact tests (see [Bhawalkar 2006]) indicate that we underestimate the amount of hardening in our model. A value of the initial hardening modulus (θ_0) that is around 0.3 of the value that we have used gives a more realistic amount of hardening. However, such a value leads to an underestimation of the flow stress at low strain rates. A better hardening model than the one used is clearly needed if we are to predict the saturation strain more accurately.

Figure 14 (left) shows the flow stress predicted by our model as a function of temperature at a plastic strain of 0.2 and for a range of strain rates. At low strain rates the flow stress decreases rapidly with temperature until it reaches the athermal value. For a strain rate of 10^3 s^{-1} and higher the initial decrease of the flow stress with temperature is small. However, as the temperature reaches the melting temperature, the flow stress drops abruptly to a small value (determined by the constitutive relation for a Newtonian

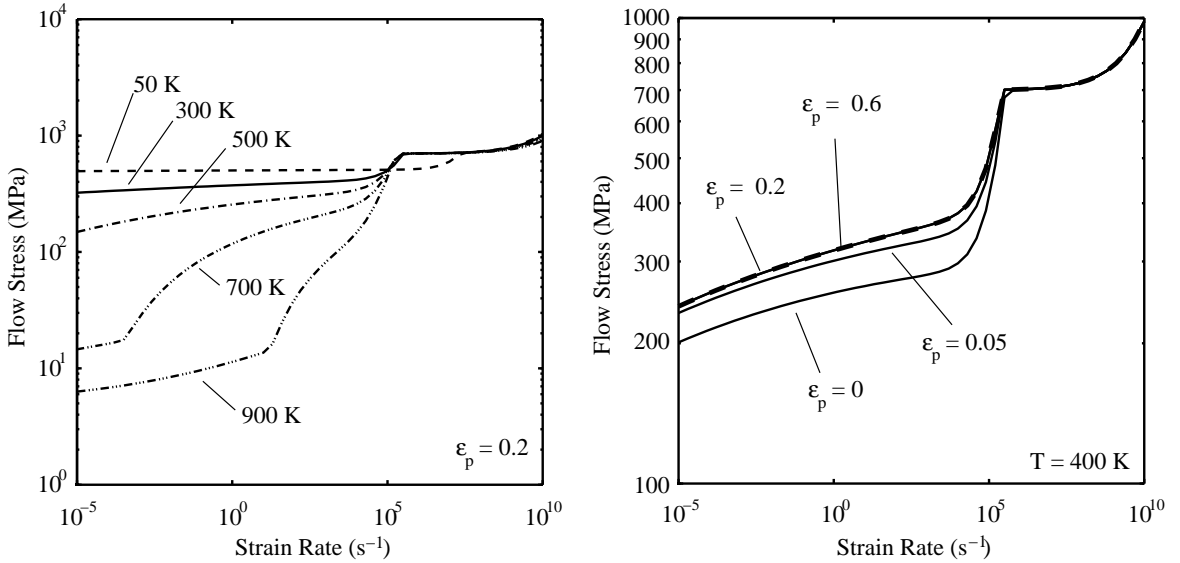


Figure 13. The predicted flow stress of 6061-T6 aluminum alloy as a function of strain rate. Left: Plastic Strain = 0.2. Right: temperature = 400 K.

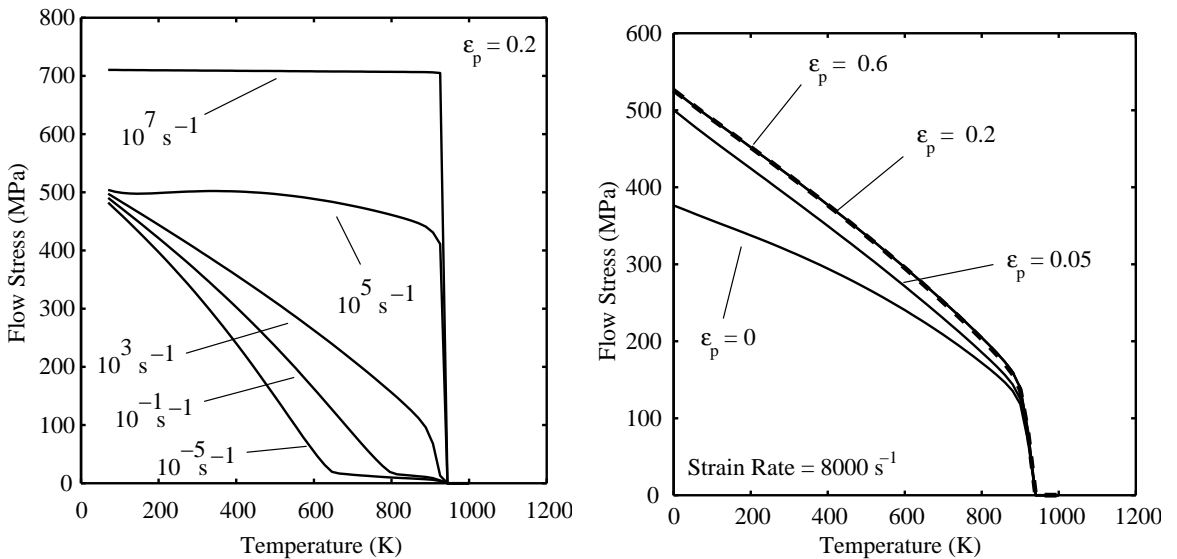


Figure 14. The predicted flow stress of 6061-T6 aluminum alloy as a function of temperature. Left: plastic Strain = 0.2. Right: strain Rate = $8000 s^{-1}$.

fluid). In the absence of experimental data, it is not obvious whether this behavior of our model accurately represents reality.

Figure 14 (right) shows the flow stress as a function of temperature at a strain rate of 8000 s^{-1} and for various levels of strain. Once again, we see that the flow stress saturates at a plastic strain of about 0.2. At low temperatures, the material hardens significantly before saturation. However, as the temperature increases, the amount of hardening decreases until at around 850 K there is essentially no hardening. This is reasonable behavior considering the fact that thermally activated dislocation motion is dominant at a strain rate of 8000 s^{-1} , and enough thermal energy is available at high temperatures to allow slip to occur more readily. The convex nature of the curves shown in the figure also indicate that we are able to capture some of the effects of high temperature solute diffusion in our model.

In Figure 15 (left) we have plotted the predicted flow stress as a function of plastic strain. The strain rate is 8000 s^{-1} and the initial temperatures range from 50 K to 900 K. Adiabatic heating is assumed, and the temperature increases with increasing strain. All the curves show a small amount of thermal softening. We also observe that the initial rate of hardening decreases with temperature. Saturation is reached at higher strains as the temperature increases. If we compare the flow stress at 300 K with those shown in Figure 12 (right), we observe that there is hardly any strain rate dependence at these strain rates. We also observe that the flow stress at 900 K is around 70 MPa which is high considering the fact that the material is close to its melting point.

Figure 15 (right) shows stress-strain plots at various strain rates. The initial temperature of the material is 400 K in these plots. Isothermal conditions were assumed for strain rates lower than 1000 s^{-1} and adiabatic conditions otherwise. Interestingly, the only curve that shows an appreciable amount of thermal softening is the curve at 1000 s^{-1} . At a strain rate of 10^5 s^{-1} , the increase in the drag coefficient with temperature offsets the amount of thermal softening that occurs. The temperature dependence of the flow

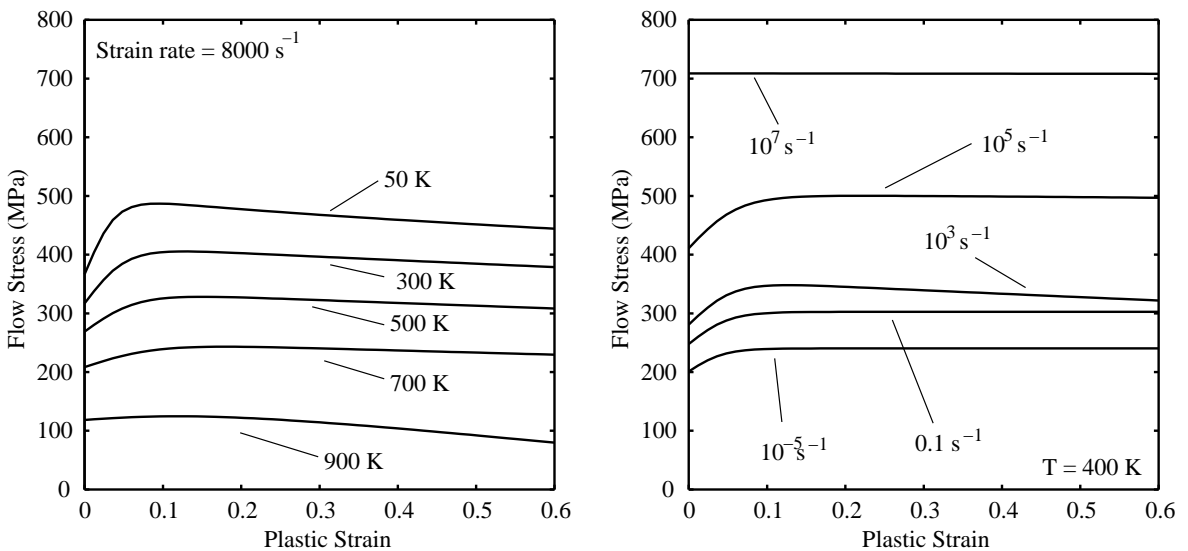


Figure 15. The predicted flow stress of 6061-T6 aluminum alloy as a function of plastic strain. Left: Strain Rate = 8000 s^{-1} . Right: temperature = 400 K.

stress at a strain rate of 10^7 s^{-1} is negligible and hence there is no appreciable thermal softening with increasing strain. The amount of strain that can be achieved at high strain rates is not clear and failure will probably occur before a plastic strain of 0.6 is achieved.

7. Concluding remarks

One of the main shortcomings of the original mechanical threshold stress model was that it was not accurate at strain rates greater than about 10^4 s^{-1} . In this paper we have presented an extended mechanical threshold stress model that is valid for strain rates at which viscous drag effects become dominant. The model has also been extended to the overdriven shock regime. The original model of Follansbee and Kocks was also designed for use in the temperature regime where solute diffusion and other thermally activated diffusion effects could be neglected (typically less than 0.6 of the melting temperature). In the extended model, we allow for higher temperatures by using a modified form of the equation for the intrinsic component of the flow stress. The strain rate dependence of the flow stress is predicted quite accurately by our model. However, the predicted temperature dependence is less accurate and could be improved. Since the density of obstacles inside a crystal decreases with increasing temperature, the activation energy should be a function of the temperature. It is possible that a temperature-dependent model for the activation energy will improve predictions of the flow stress of 6061-T6 aluminum at high temperatures. More high temperature–high strain rate data of the type provided by [Rosenberg et al. \[1986\]](#) are essential before such a model can be developed.

We have provided a detailed description of the procedure for calculating the drag stress. Two methods for computing the viscous drag coefficient have been compared in the paper. We have found that Liebfried's theory provides an estimate of the viscous drag coefficient that better matches experiment. We have also added a temperature dependence to the dislocation evolution equations of Estrin and Kubin and determined new parameters for these equations that fit experimental data. Estrin and Kubin have estimated that the mobile dislocation density is around 10^{14} m^{-2} at saturation. Our results suggest that a more accurate estimate of this quantity is $2 \times 10^{11} \text{ m}^{-2}$.

The evolution equations of the internal variable (mechanical threshold stress) have been simplified and a nondimensional form for the initial hardening modulus has been proposed. The simplified equations allow for an exact expression for the internal variable as a function of plastic strain which can be evaluated efficiently in a computational code. Alternatively, we could have used the equations for the evolution of the dislocation density with the dislocation density as an internal variable. However, the system of equations involved is very stiff and computationally expensive to solve for high strain rate problems. The parameters for 6061-T6 aluminum alloy (needed for the evolution of the internal variable) that we have determined in this paper lead to an underestimation of the saturation stress and the amount of hardening. This effect is particularly noticeable for high hydrostatic pressures. Also, the pressure dependence of the flow stress is not reflected completely by the pressure dependence of the shear modulus (for which a model is presented in the paper). Explicitly pressure–dependent models for the hardening modulus and the saturation stress are clearly needed.

In addition, we have developed an equation for the evolution of temperature with plastic strain based on thermodynamic principles. An expression has been presented for the Taylor–Quinney coefficient as a function of the mechanical threshold stress internal variable. This expression requires a functional

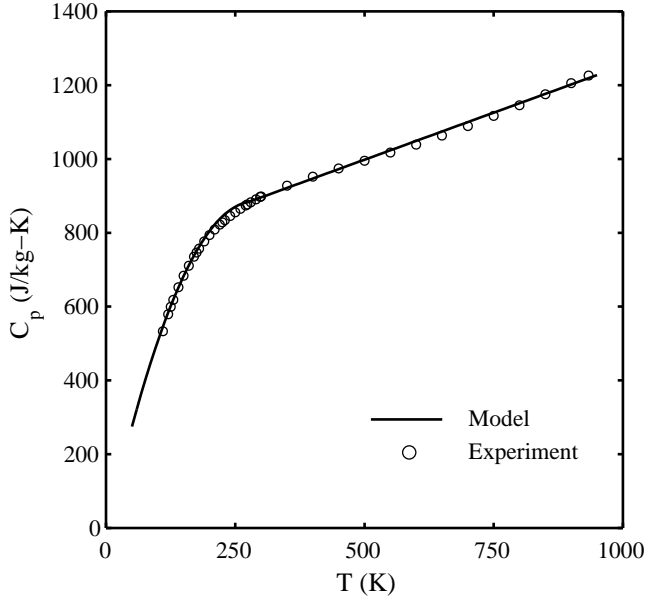


Figure 16. Specific heat of aluminum. The experimental data, shown as open circles, are from [Desai 1987]. The solid line shows the prediction of the model in Equation (31).

representation of the Gibbs free energy density. We have postulated a form of the derivatives of this function that matches experimental data qualitatively. However, further research is needed to arrive at a better form of the Gibbs free energy density function.

Important issues that have not been addressed in this paper are cyclic loading and kinematic hardening models, deformation induced anisotropy in both the elastic and plastic behavior, the appropriate form of the yield condition and the flow rule, and the modeling of void nucleation and growth and the accompanying elastic and plastic damage. Also, in this paper, we have considered only one dimensional analytical verification tests. Consideration of some of the above issues and detailed numerical validation experiments using the material point method Sulsky et al. [1994; 1995] are in progress and will be the subject of a subsequent paper.

Appendix A: specific heat, EOS, melting temperature, shear modulus

Experimental data indicate that the specific heat (at constant pressure) of aluminum increases by around 37% between room temperature and melting [Desai 1987]. A least squares curve fit to the experimental data yields the following parameters for the model given in Equation (31):

$$\begin{aligned} A_1 &= -11.73 \text{ J/kg-K}, & B_1 &= 6.28 \text{ J/kg-K}^2, \\ C_1 &= -0.011 \text{ J/kg-K}^3, & A_2 &= 743.13 \text{ J/kg-K}, \\ & & B_2 &= 0.51 \text{ J/kg-K}^2. \end{aligned}$$

A plot of the fit is shown in Figure 16.

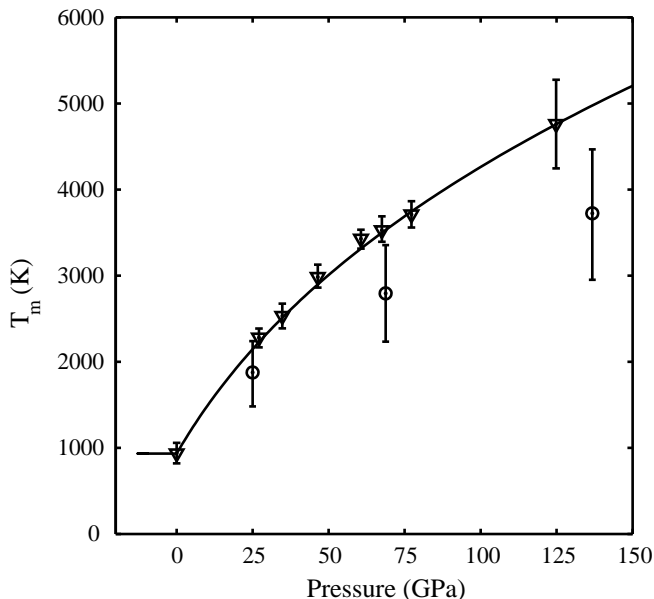


Figure 17. Melting temperature of aluminum. The triangles show experimental data from [Boehler and Ross 1997] and the circles show experimental data collected in [Burakovsky et al. 2000]. The solid line shows the prediction of the model in Equation (29).

The shear modulus model given in Equation (26) requires a pressure. We use the equation of state (EOS) in Equation (30) to compute the pressure for a given change in density. The parameters that we have used for 6061-T6 aluminum are $B_0 = 76.3$ GPa and $B'_0 = 4.29$ [Burakovsky and Preston 2005]. Experimental data for 6061-T6 aluminum alloy [Vaidya and C. 1970], 2024-O aluminum alloy [McQueen et al. 1970], and pure aluminum [Mitchell and Nellis 1981] are closely matched by the EOS model up to a compression ratio (ρ/ρ_0) of 1.4.

Though the alloying content can have a significant effect of the melting temperature of metals, we have assumed in this paper that the melting temperature of 6061-T6 aluminum does not differ significantly from that of pure aluminum. The parameters of the model of melting temperature given in Equation (29) for 6061-T6 aluminum are taken to be

$$T_{m0} = 933.6 \text{ K}, \quad q = 3.3,$$

$$\Gamma_1 = 0.84(\text{gm/cc})^{1/3}, \quad \Gamma_2 = 45.4(\text{gm/cc})^q.$$

We also assume that for hydrostatic tensile states the melting temperature is equal to T_{m0} . The prediction of the model of Equation (29) is compared with experimental data in Figure 17. An initial density of 2700 kg/m^3 was used while computing the quantities in the figure. The model fits the data from [Boehler and Ross 1997] quite well (triangles) but not the older data collected in [Burakovsky et al. 2000] (circles). The accuracy of the Boehler and Ross data has been confirmed by more recent numerical experiments by Voadlo and Alfe [2002].

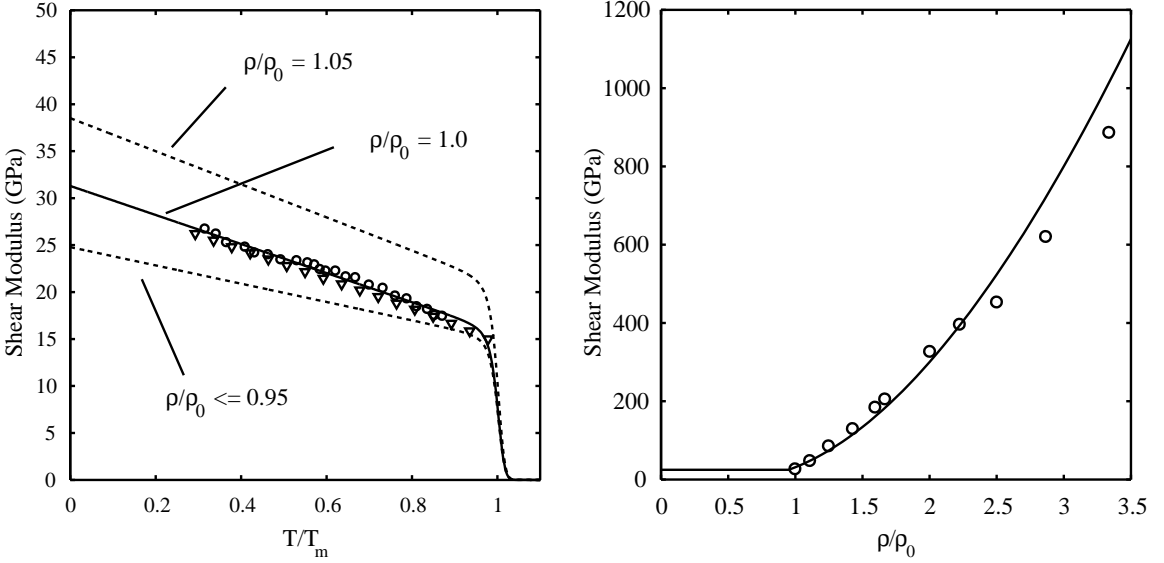


Figure 18. Shear modulus of 6061-T6 aluminum alloy. Left: shear modulus vs. temperature. Right: shear modulus vs. mass density at 0 K. At left, the triangles show experimental data for pure aluminum from [Tallon and Wolfenden 1979] and the circles show experimental data for 6061-T6 aluminum alloy from [Brown et al. 1996]. The solid line shows the prediction of the model in Equation (26) at ambient pressure. The dotted lines show predicted shear moduli under hydrostatic tension and compression. At right, the circles show results from electronic structure calculations [Burakovsky et al. 2003] and the solid line shows the prediction of our model.

The shear modulus model in Equation (26) allows for a smooth transition to zero shear modulus beyond melting. This feature is advantageous in numerical simulations for conditions where melting might occur. The parameters of the shear modulus model for 6061-T6 aluminum are $\mu_0 = 31.3$ GPa, $\partial\mu/\partial p = 1.8$ [Guinan and Steinberg 1974], $a_1 = 0.53$, $a_2 = 0.10$, $a_3 = 0.37$ [Burakovsky and Preston 2005], $C = 0.049$ [Nadal and Le Poac 2003], $\zeta = 0.1$, and $m = 26.98$ amu. The predicted shear modulus as a function of temperature is compared with experimental data in Figure 18 (left). An initial density of 2700 kg/m^3 was used for the calculations in the figure. Further details and comparisons with other models can be found in [Bhawalkar 2006].

The density dependence of the shear modulus at 0 K is compared with computed results in Figure 18 (right). Since the pressure depends directly on the density in the equation of state, this plot may also be considered to be a measure of the pressure dependence of the shear modulus. Our model predicts the shear modulus quite accurately up to $\rho/\rho_0 = 2.2$. Beyond this point the material undergoes a phase transition to the bcc phase and our model overestimates the shear modulus with increasing pressure.

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