THERMOMECHANICAL FATIGUE MADE EASY

Darrell Socie^{*} and Benjamin Socie⁺

Thermo Mechanical Fatigue (TMF) is a complex subject because of the interaction of several failure modes including fatigue, oxidation and creep with a wide variety of complex thermal and mechanical loads. Many models exist and several commercial software codes are available for evaluating the durability of a structure under these conditions. These codes often require the user to be a fatigue specialist with considerable training. In addition, the material properties needed for these calculations are difficult and very expensive to obtain experimentally. In this paper an approximate procedures to obtain estimates of the material properties and a simple software system for the non-specialist are described.

INTRODUCTION TO THERMOMECHANICAL FATIGUE

Thermomechanical fatigue (TMF) is caused by combined thermal and mechanical loading where both the stresses and temperatures vary with time. This type of loading can be more damaging by more than an order of magnitude compared with isothermal fatigue at the maximum operating temperature. Material properties, mechanical strain range, strain rate, temperature, and the phasing between temperature and mechanical strain all play a role in the type of damage formed in the material. These types of loadings are most frequently found in start-up and shut-down cycles of high temperature components and equipment. Typically design lives are a few thousand cycles and involve significant plastic strains.

One of the major differences between isothermal and thermal mechanical fatigue is constraint. When heated, structures develop thermal gradients as they expand. Expansion near stress concentrators is often constrained by the surrounding cooler material. In this case thermal strain is converted into mechanical strain which causes fatigue damage in the structure. Total constraint exists when all of the thermal strain is converted into mechanical strain. Over constraint can occur in a stress concentration where the mechanical strain is greater than the thermal strain. One measure of the degree of constraint is the ratio of the thermal and mechanical strain rates.

TMF loading is often described to be in-phase (IP) or out-of-phase (OP). A schematic illustration of the stress-strain response under these two loadings is given in Fig. 1. In IP loading the maximum temperature and strain occur at the same time. In OP loading, the material experiences compression at highest temperature and tension at lower temperatures. IP loading is more likely to cause creep damage during tensile stresses at high temperatures. OP loading is more likely to cause oxidation damage because an oxide film can form in compression at the higher temperature and then rupture during the subsequent low temperature tensile portion of the loading cycle where the oxide film is more brittle.

*Mechanical Science and Engineering, University of Illinois at Urbana-Champaign, Urbana Illinois, USA

⁺ BBM Plus, Saint Joseph Illinois, USA

Figure 2 shows early TMF test data from Jaske[1] for a low carbon steel. Isothermal fatigue tests at various temperatures are shown as the lines without symbols. TMF OP tests are shown with the solid symbols in the figure. Fatigue lives for tests conducted with variable strains and temperatures between 93°C and 538°C are more than an order of magnitude shorter than tests conducted at a constant temperature of 538°C. Assuming the "worst case" isothermal material properties is very non-conservative and a TMF fatigue analysis must be employed for TMF assessments.

There are many active mechanisms in the TMF process. For discussion it is convenient to consider damage from three primary sources: fatigue, oxidation and creep[2]. Damage from each process is summed to obtain an estimate of the total fatigue life, N_f .

$$\frac{1}{N_{f}} = \frac{1}{N_{f}^{fatigue}} + \frac{1}{N_{f}^{oxidation}} + \frac{1}{N_{f}^{creep}}$$
(1)

Frequently one of the damage mechanisms is dominant. From a modeling prospective, this suggests that the individual damage models and their associated material properties must be accurate only for those conditions where the life is dominated by that failure mechanism.

Fatigue mechanisms

Fatigue damage in the life regime of interest in TMF is in the form of nucleation and growth of microcracks. Figure 3 shows an example of the surface damage observed at higher plastic strains[3]. Microcrack density will decrease at lower strain levels. Many microcracks nucleate on the surface and a few of them are able to penetrate into the bulk of the material. The process is driven by cyclic plastic strains where oxidation and creep effects are negligible. Fatigue damage will dominate at high strain ranges, strain rates and low temperatures.

Oxidation mechanisms

Oxidation damage can occur in the form of an oxide intrusion such as the one shown in Fig. 4 [4]. In OP loading, an oxide layer can form on the surface when the material is hot and in compression. At the lower temperature the oxide layer becomes brittle. During mechanical straining it then cracks to expose new clean metal surfaces. This clean metal will rapidly oxidize and the process repeats during the next mechanical strain cycle. Ultimately this will form a crack which can then grow during the mechanical strain cycle. Oxide cracks can also form during IP loading. In this case, the oxide forms during the hot portion of the loading cycle while the material is in tension. Then upon cooling the oxide film undergoes a buckling delamination which fractures the oxide and exposes clean metal surfaces. Oxide formation and rupture during isothermal loading is not the dominant failure mechanism and is not reflected in isothermal test or materials data. Oxide formation will occur easier and faster at higher temperatures. The tendency of the oxide to develop microcracks will depend on the cyclic strain range, stress does not play a role in the development of oxide induced microcracks.

Creep mechanisms

Creep is essentially a diffusion process. At high stresses diffusion allows dislocations to climb over barriers. At lower stresses diffusion occurs along the grain boundaries. Figure 5 shows a triple point crack that has formed at the junction between grains[5]. These microcracks form as a result of grain boundary sliding to accommodate the change in shape of elongated grains. Diffusion is highly temperature and time dependant. Maximum stress rather than strain range has a dominant role in the formation of these microcracks. The interaction of the strain rate and temperature has a strong influence on the stresses that are observed during cyclic loading.

One common feature of all these mechanisms is that they involve the nucleation and early growth of microcracks. There will also be interactions between the mechanisms as well. For modeling, individual components are considered and interactions are ignored. This notation is employed in the following equations: cr - creep, in – inelastic, mech – mechanical, ox - oxidation, and th – thermal. The traditional nomenclature is employed for stress, σ , strain, ϵ , strain rate, $\dot{\epsilon}$, temperature, T, and life N.

Fatigue Damage Model

Conventional low cycle fatigue damage is a surface phenomena where small microcracks nucleate and grow on the surface of the material. Bulk stresses and strains are employed to describe fatigue damage because the microcracks growth is too complex to describe in detail. The strain-life equation is the most common description of the process.

$$\frac{\Delta \varepsilon}{2} = \frac{\sigma_{f}}{E} \left(2N_{f}^{\text{fatigue}} \right)^{b} + \varepsilon_{f}^{'} \left(2N_{f}^{\text{fatigue}} \right)^{c}$$
(2)

- σ_{f} fatigue strength coefficient
- b fatigue strength exponent
- ε_{f} fatigue ductility coefficient
- c fatigue ductility exponent
- E elastic modulus

Oxidation Damage Model

The oxidation damage formulation of Neu and Schitoglu[6] is employed in this study. Oxide damage will occur when the strain range exceeds a threshold for oxide cracking.

if
$$\Delta \varepsilon_{\text{mech}} > \varepsilon_0$$
 $\frac{1}{N_{\text{f}}^{\text{oxidation}}} = \left[\frac{H_{\text{cr}}}{\Phi^{\text{ox}} K_{\text{peff}}}\right]^{-\frac{1}{\beta}} \frac{2(\Delta \varepsilon_{\text{mech}})^{\frac{2}{\beta}+1}}{\dot{\varepsilon}^{1-\frac{5}{\beta}}}$ (3)

 ϵ_{o} threshold strain for oxide cracking

H_{cr} constant related to critical oxide thickness

- β mechanical strain range exponent
- b thermal strain rate sensitivity exponent

$$\Phi_{\rm ox} = \frac{1}{t_{\rm c}} \int_{0}^{t_{\rm c}} \phi_{\rm ox} dt \tag{4}$$

$$\phi_{\text{ox}} = \exp\left[-\frac{1}{2}\left(\frac{\dot{\epsilon}_{\text{th}} / \dot{\epsilon}_{\text{mech}} + 1}{\xi^{\text{ox}}}\right)^2\right]$$
(5)

 ξ^{ox} oxidation phasing constant for thermal and mechanical strains

$$K_{peff} = \int_{0}^{t_{c}} D_{0} \exp\left(\frac{-\Delta H^{ox}}{RT}\right) dt$$
(6)

 ΔH^{ox} activation energy for oxidation

D_o scaling constant for oxidation

Oxidation damage is a function of the strain range, strain rate, and temperature. A phasing factor ϕ_{ox} is introduced to account for the type of oxide cracking that occurs in either IP or OP loading. Phasing is represented by the ratio of thermal and mechanical strain rates. Oxidation rate is determined by the effective parabolic oxidation constant, K_{peff} .

Creep Damage

The creep damage formulation suggested by Neu and Schitoglu[6] is also employed in this study.

$$\frac{1}{N_{f}^{creep}} = \int_{0}^{t_{c}} A_{cr} \Phi^{cr} \exp\left(\frac{-\Delta H^{cr}}{RT}\right) \left(\frac{\alpha_{1}\overline{\sigma} + \alpha_{2}\sigma_{h}}{K}\right)^{m}$$
(7)

- ΔH^{cr} activation energy for creep
- A_{cr} scaling constant for creep
- m creep stress exponent
- α_1 stress state constant
- α_2 hydrostatic stress sensitivity constant

$$\Phi_{\rm cr} = \frac{1}{t_{\rm c}} \int_{0}^{t_{\rm c}} \phi^{\rm cr} dt$$
(8)

$$\phi_{\rm cr} = \exp\left[-\frac{1}{2}\left(\frac{\dot{\varepsilon}_{\rm th} / \dot{\varepsilon}_{\rm mech} - 1}{\xi^{\rm cr}}\right)^2\right]$$
(9)

 ξ^{cr} creep phasing constant for thermal and mechanical strains

Creep damage is a function of the stresses, time and temperature. Microstructural creep damage differs in tension and compression. It is commonly assumed that microcracks do not form and grow

in compression. If no creep damage occurs in compression $\alpha_1 = 1/3$ and $\alpha_2 = 1$. Here, K is the drag stress which will be defined in the next section. A phasing factor ϕ_{cr} is also introduced to account for different creep damage mechanisms such as intergranular or transgranular cracking.

Constitutive Equation Model

A unified constitutive model first suggested by Bodner and Partom[7] is employed to compute the stresses. The combined effects of both creep and plasticity are treated as inelastic strains. At lower stresses, time dependant creep dominates the behavior. Plasticity dominates at higher stresses. A drag stress, K, is introduced into the formulation. The drag stress is an internal state variable that is related to the strength of the material. It is the stress that defines the transition from creep to plasticity dominated deformation. It is not constant but depends on the temperature.

$$\dot{\epsilon}^{\text{in}} = \begin{cases} \frac{A_0 \left(\frac{\overline{\sigma}}{K}\right)^{n_1} \exp\left(\frac{-\Delta H^{\text{in}}}{RT}\right) & \left(\frac{\overline{\sigma}}{K}\right) \le 1\\ \frac{A_0 \exp\left[\left(\frac{\overline{\sigma}}{K}\right)^{n_2} - 1\right] \exp\left(\frac{-\Delta H^{\text{in}}}{RT}\right) & \left(\frac{\overline{\sigma}}{K}\right) \ge 1 \end{cases}$$
(10)

- ΔH^{in} activation energy for inelastic deformation
- A_o scaling constant for inelastic deformation
- n₁ exponent for creep dominated deformation
- n₂ exponent for plasticity dominated deformation

A linear temperature for the drag stress is often employed. Other forms are possible.

$$\mathbf{K} = \mathbf{K}_0 - \mathbf{K}_1 \mathbf{T}$$

- K₀ back stress
- K₁ back stress temperature dependence

A linear dependence for the elastic modulus is also frequently employed.

$$E = E_0 - E_1 T$$

- E₀ elastic modulus
- E₁ elastic modulus temperature dependence

The thermal expansion coefficient is also needed in the analysis to determine thermal strains.

 α thermal expansion coefficient

These TMF equations represent a model for steady state deformation and require a total of 27 material modeling constants.

SIMPLIFIED MATERIAL PROPERTIES

TMF mechanisms are complicated and influenced by the material microstructure, environment and external loading and a complete set of material data is preferred. Twenty seven material modeling constants means that only a few materials have been fully characterized. Yet, life assessments must frequently be made in the early stages of design before a complete set of materials data is available. As a result, there is a need to make estimates of the material properties from other more readily available data. In this paper, simplified material properties are employed. They are based on a classification system, low carbon steels, alloy steels, aluminum etc. because a complete set of data is available for only a few materials. These materials will be designated as reference materials. Properties from these materials will be modified to account for microstructural differences between materials within a given class. Fatigue constants are always needed for the analysis.

Both creep and oxidation damage models have a phasing factor. The phasing factor is shown in Fig. 6 as a function of the thermal and mechanical strain ratio. This factor determines the dominant failure mechanism, creep or oxidation. Creep will dominate IP TMF loading. Oxidation will dominate for both isothermal and OP TMF. It should be noted that many TMF problems involve constrained heating and cooling which result in out-of-phase loading. In this case, knowledge of the creep properties are unnecessary because only the oxidation and fatigue behavior is important.

Fatigue Constants

Many correlations between fatigue and tensile properties have been proposed. Many of them have been validated only for steels. Muralidharan and Manson's[8] method has been validated for wide range of materials. It is based on the elastic modulus, E, ultimate strength, S_u , and true fracture strain, ε_f .

$$\frac{\Delta\epsilon}{2} = 0.623 \left(\frac{S_u}{E}\right)^{0.832} (2N_f)^{-0.09} + 0.0196 (\epsilon_f)^{0.155} \left(\frac{S_u}{E}\right)^{-0.53} (2N_f)^{-0.56}$$
(11)

Oxidation Constants

Oxidation is dominated by the matrix material rather than microstructure in most alloys. As a first approximation, alloys of a similar matrix are expected have the same oxidation behavior. No adjustments are needed for microstructure and all materials of the same class will have the same behavior.

Creep Constants

Creep is a process that is driven by diffusion either in the bulk material or along the grain boundaries. Creep damage should be directly related to the creep rate and rupture life. Equation 7 for creep damage can be divided into two terms, one for temperature and one for stress dependence. Figure 7 shows a Larson-Miller plot for various alloys. Two materials with the same value of the

Larson-Miller parameter, P_{LM} , will have the same time and temperature dependence. Note that the lines describing the material behavior are nearly parallel for a given class of materials. For example, the difference between low carbon and Cr-Mo steel is essentially a shift in stress level. This observation allows us estimate the behavior of other materials in the same class. The activation energy for creep, ΔH^{cr} , is directly related to the self-diffusion activation energy which will depend on the matrix material not the microstructure. The stress constants, α_1 and α_2 , and the phasing constant, ξ^{cr} , can also be considered functions of the matrix material and as a first approximation will not change with the microstructure. Exponents such as m are an indication of the mechanism and are not expected to change. This leaves a single constant, A_{cr} , that will depend on the microstructure. As a first approximation, this constant can be scaled from the reference material data. Two materials with the same P_{LM} will have the same creep damage and the integrand of Eq. 7 must also be the same for both materials. In Fig. 7 consider low carbon and C-Mo steel. The allowable stresses for C-Mo steel are shifted up by about 25%. Let the creep strength be denoted as S and the creep strength of the reference material as $S|_{ref}$. Manipulating Eq. 7 and eliminating the temperature terms results in an approximate expression for the constant A_{cr} .

$$A_{cr} = A_{cr}|_{ref} \left(\frac{\frac{S_{ref}}{S}}{\frac{K_{ref}}{K}} \right)^{m}$$
(12)

Microstructural effects are indirectly included in the creep damage formulation by normalizing the stress by the drag stress K. Drag stress will also be directly related to the materials creep strength since the drag stress represents the transition between creep and plasticity dominated behavior. Then Eq. 12 can be further simplified to $A_{cr} = A_{cr}|_{ref}$. The final result is that no adjustments are needed for the creep constants. Differences between materials within a class are modeled by changes in the drag stress.

. .

Constitutive Equation Constants

Stresses are not included in the fatigue and oxidation models, only the creep damage model. In the unified constitutive models no attempt is made to separate the creep and plasticity strains, they are all considered as inelastic strains. The exponential terms in Eq. 10 are related to high temperature creep deformation. Following the same arguments used for determining creep damage constants, the activation energy and exponents are expected to be a constant for alloys within a class of materials. The remaining two constants A_0 and K will depend on the material microstructure. Drag stress should scale directly with the materials strength, either creep strength or yield strength. The constant A_0 is assumed to remain constant.

A very simple model for approximating TMF material constants is proposed, simply adjust the material drag stress in proportion to the material strength. Creep strength is preferred but room temperature yield strength could be employed when creep data is not available.

FATIGUECALCULATOR

To encourage more widespread use of fatigue analysis, a website <u>www.fatiguecalculator.com</u> has been developed to perform routine fatigue damage calculations. The website is free and open to everyone to use. TMF analysis is included with an appropriate material database of TMF constants for various materials. The TMF portion consists of three sections, definition of the duty cycle, material properties and results. Figure 8 shows an example of the duty cycle input. It is in the form of a list of endpoints for each loading segment in the duty cycle for the strain, time and temperature. The loading may be specified as mechanical strains, total strains or stress or strain hold times. The materials database includes several materials and the user can enter their own properties. Results of the analysis are shown in Fig. 9. In addition to the computed fatigue lives, several plots of stresses and strains are provided.

COMPARISON WITH EXPERIMENTS

Two of the datasets analyzed are presented here. The basis of comparison is the computed TMF lives because our objective is to make estimates of fatigue lives not reproduce material modeling parameters. The first dataset analyzed is that shown in Fig. 2 for 1010 steel. The reference data employed was 1070 steel[6]. Creep strength data was not available for both materials so the yield strength was used. Fatigue constants were obtained from the SAE handbook[10] for 1010 steel. Yield strengths for 1010 and 1070 steel were obtained from www.matweb.com. It lists the yield strengths as 300 and 580 MPa for 1010 and 1070 steel respectively. Before performing the calculations the drag stress constants were reduced by a factor of 0.52. No other changes in the oxidation, creep or constitutive equation constants were made. Results of the analysis are given in Table 1. In addition to the experimental and calculated fatigue lives, the relative contribution of each of the three damage mechanisms is given.

Mechanical T _{max}		Experiment Analysis Cycles Cycles		Fatigue Damage	Oxidation Damage	Creep Damage	
OP	0.0200	538	116	427	60%	40%	-
	0.0100	538	157	1850	51%	49%	-
	0.0068	538	262	4290	43%	57%	-
	0.0041	538	632	13400	30%	70%	-
	0.0148	427	600	1330	94%	6%	-
	0.0071	427	1307	8350	90%	10%	-
	0.0040	427	4890	40200	83%	17%	-
	0.0143	316	569	1530	99%	1%	-
	0.0070	316	3232	9240	99%	1%	-
IP	0.0200	538	75	386	55%	32%	13%
	0.0100	538	489	836	23%	18%	59%
	0.0070	427	3282	7870	85%	8%	7%
	0.0072	317	4778	8570	100%	-	-

TABLE 1 1010 Steel results from Jaske [1]

A second set of test data [11] for a Ni based superalloy IN 738LC was also analyzed. The reference material selected was another Ni based superalloy Mar M247[12]. Creep strength of IN

738 was taken from [13] as 210 MPa at 870 C. Similar data for Mar M247 was obtained from [14] as 280 MPa. Again the drag stress constants for Mar M247 were reduced by 0.75 to obtain an estimate of the IN 738 constants. No other adjustments were made to the Mar M247 oxidation, creep or constitutive equation constants were made. Strain life data was taken from reference 11. Results of the analysis are shown in Table 2.

s S	Mechanical strain Range	Experiment Cycles	Analysis Cycles	Fatigue Damage	Oxidation Damage	Creep Damage
OP	0.008	100	182	9%	91%	-
	0.004	4000	1620	1%	99%	-
IP	0.0075 0.005 0.004	180 1000 2000	77 837 2490	3% 2% 1%	- -	97% 98% 99%

TABLE 2 IN 738LC results from Fleury and Ha [11]

Results in Tables 1 and 2 show that the simple approximate method can be expected to produce fatigue lives that are within an order of magnitude of the experimental lives.

SUMMARY

A simple method for determining material modeling constants for TMF damage calculations has been proposed. The method consists of using reference materials for which the constants are known and then adjusting the drag stress to obtain constants for other similar materials. A web based analysis procedure was also described.

REFERENCE LIST

- (1) Jaske, C.E. Thermal Fatigue of Materials and Components, ASTM STP 612, 1976, pp. 170-198
- (2) Schitoglu, H. Advances in Fatigue lifetime Predictive Techniques, ASTM STP 1122, 1992, pp. 47-76
- (3) Stolarz, J, Ecole Nationale Superieure des Mines, Presented at LCF 5 in Berlin, 2003
- (4) Viswanathan, R. and J. Stringer, *J. of Engineering Materials and Technology*, Vol. 122, July 2000, pp. 246-255
- (5) Morris, D.G. and D.R. Harries, J. of Materials Science, Vol. 12, No. 8, 1977, pp. 1587-1597
- (6) Neu, R.W. and H Sehitoglu, *Metallurgical Transactions A*, Vol. 20A, 1989, pp. 1769-1783
- (7) Bodner, S.R. and Y. Partom, J. of Applied Mechanics, Vol. 46, 1979, pp. 805-810

- (8) Muralidharan, U and S.S. Manson, J. Engineering Materials and Technology, Vol. 110, 1988, pp. 55-58
- (9) Dowling, N.E. Mechanical Behavior of Materials, Prentice Hall, 1998
- (10) SAE Handbook, Society of Automotive Engineers, 2003
- (11) Fleury E. and J.S. Ha Materials Science and Technology, Vol. 17, 2001, pp. 1087-1091
- (12) Sehitoglu, H. and D.A. Boismier, J. Engineering Materials and Technology, Vol. 112, 1990, pp.80-89
- (13) *High Temperature High Strength Nickel Base Alloys*, International Nickel Company, 3rd Edition, 1977
- (14) Superalloys A Technical Guide, ASM International, 2nd Edition, 2002







Figure 2 TMF Test Results for 1010 Steel [1]



surface view

cross section view

FIGURE 3 Formation of Surface Cracks in 20-25 Austenitic Steel [3]



Figure 4 Oxidation Damage in Steel [4]



Figure 5 Wedge crack nucleation in Type 316 stainless steel [5]



Figure 7 Larson-Miller Plot [9]

Loading

bottom of

You may enter the loading as a short, 10 points or less, load history or as a sine wave. To enter a load history, enter values into the first set of boxes. To enter parameters in the second set of boxes. The Plot button below is for you to verify that the correct loading information was entered.



Figure 8 TMF Duty Cycle Definition



