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Cyclic Thermo-Mechanical Material Modelling And Testing Of 316 Stainless Steel

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Abstract

A programme of cyclic mechanical testing of a 316 stainless steel, at temperatures of up to 600°C under isothermal conditions, for the identification of material constitutive constants, has been carried out using a thermo-mechanical fatigue test machine (with induction coil heating). The constitutive model adopted is a modified Chaboche unified viscoplasticity model, which can deal with both cyclic effects, such as combined isotropic and kinematic hardening, and rate-dependent effects, associated with viscoplasticity. The characterisation of 316 stainless steel is presented and compared to results from tests consisting of cyclic isothermal, as well as in-phase and out-of-phase thermo-mechanical fatigue conditions, using interpolation between the isothermal material constants to predict the material behaviour under anisothermal conditions.

Keywords: Thermo-mechanical fatigue, Creep, Plasticity, Chaboche unified viscoplasticity model, 316 stainless steel

1. Introduction

Thermo-mechanical fatigue, or TMF, has received an increasing amount of attention over the last thirty years. The two most common experimental TMF waveforms are in-phase (IP) and out-of-phase (OP) as shown in Figure 1, where $\phi$ represents the phase angle between the strain and temperature waveforms. Some of the areas of particular interest have included the effect of phase angle on the life and failure of materials undergoing TMF as shown by Pahlavanyali et al [1], TMF behaviour of superalloys, e.g. Evans et al [2], TMF of specific components, e.g. Harrison et al [3], and crack growth under TMF conditions, e.g. Dai et al [4].
Unified material models are considered as a robust way of modelling the behaviour of materials where both rate-independent and rate-dependent effects occur simultaneously, e.g. for high temperature cyclic loading. The Chaboche unified viscoplasticity model, for example, includes both non-linear isotropic hardening and kinematic hardening, for cyclic phenomena such as hardening and the Bauschinger effect [5], as well as creep effects. This model was first published by Chaboche and Rousselier in 1983 [6, 7] and is discussed further by LeMaitre and Chaboche [8]. Since then the model has been widely used, for example, Tong and Zhan et al [9-14].

The present paper is particularly concerned with the application of the Chaboche model to anisothermal conditions. In order to achieve this, isothermal tests have been performed to obtain the material constants at a range of temperatures. At each temperature, the model has been used to simulate isothermal cyclic conditions and the predictions compared to the experimental data. Each material constant was then represented as a function of temperature for simulation of the anisothermal IP and OP TMF conditions. The results from these simulations are compared to experimental TMF data. Examples of previous work on TMF of stainless steel includes that of Santacreu et al [15] concerned with automotive exhaust applications and Rau et al [16] in the exploration of more complex testing conditions, to understand more realistic behaviours of materials in industrial use.

2. Experimental program

All of the results presented have been obtained using 316 stainless steel specimens. Table 1 shows the chemical composition of the 316 stainless steel obtained as an average of three spark emission tests performed on material from the same batch as the specimens.

<table>
<thead>
<tr>
<th>( \text{Fe} )</th>
<th>( \text{Cr} )</th>
<th>( \text{Ni} )</th>
<th>( \text{Mo} )</th>
<th>( \text{Mn} )</th>
<th>( \text{Si} )</th>
<th>( \text{Cu} )</th>
<th>( \text{V} )</th>
<th>( \text{Co} )</th>
<th>( \text{S} )</th>
<th>( \text{C} )</th>
<th>( \text{Nb} )</th>
<th>( \text{P} )</th>
<th>( \text{W} )</th>
<th>( \text{Ti} )</th>
<th>( \text{Al} )</th>
</tr>
</thead>
<tbody>
<tr>
<td>66.4</td>
<td>16.8</td>
<td>11.8</td>
<td>2.15</td>
<td>1.42</td>
<td>0.5</td>
<td>0.49</td>
<td>0.08</td>
<td>0.07</td>
<td>0.03</td>
<td>0.02</td>
<td>0.02</td>
<td>&lt;0.02</td>
<td>0.01</td>
<td>0.0</td>
<td>0.01</td>
</tr>
</tbody>
</table>

2.1. TMF machine and thermal calibration

The TMF machine used in this work is an Instron 8862 thermo-mechanical fatigue system which utilises radio-frequency (RF) induction heating and forced air cooling through the centre of the specimen in order to achieve rapid heating and cooling, respectively. The maximum achievable load from the machine is 35kN (limited by the grips) and the maximum allowable temperature is 1100°C.
The requirement for temperature uniformity in the gauge section of the specimen during testing was that the entire gauge section was within ±10°C of the target temperature. Therefore initial experimental work was concerned with achieving this.

Thermocouples were placed along the gauge section of the specimen in order to monitor the axial, as well as the circumferential temperature gradients within the specimen gauge section during a series of ramp and hold, as well as cyclic thermal testing. Initially the results were not within this tolerance, with axial deviations of up to ±30°C from the target temperature. Therefore new coil designs were investigated.

![Photograph of the heated specimen, induction coil and extensometer setup on the TMF machine.](image)

A key problem faced was achieving the temperature uniformity required whilst leaving enough space between the turns of the coil for the attachment of the extensometer to the gauge section of the specimen. Figure 2 shows the final coil design which gives temperature uniformity throughout the gauge section to within the tolerance required for target temperatures ranging from 200°C to 1000°C. Figure 3 shows the temperature uniformity results obtained using this coil and a 316 stainless steel specimen for a target temperature of 800°C.

![Thermocouple positions and thermal uniformity results using the final coil.](image)

2.2. Isothermal cyclic testing

Isothermal cyclic tests were carried out at temperatures of 300°C, 500°C, 550°C and 600°C. Figure 4 shows the specimen geometry used in these tests. At each temperature, the test was performed for fifty loops at four strain ranges, i.e. stepped strain-range testing (200 loops in total),
using one specimen for each temperature. Example results at 600°C from these tests are shown in Figure 5. It can be seen that the majority of the material hardening occurred at the first and lowest strain range (±0.3%). Therefore these test results at a strain range of ±0.3% were used to obtain the material constants for the Chaboche unified viscoplasticity model (see Sections 3 and 4) for each temperature.

Figure 4. Specimen geometry used on the TMF machine.

Figure 5. Isothermal cyclic test results at 600°C.

2.3. TMF

The results of two TMF tests, carried out under IP and OP conditions are presented. The IP TMF test was carried out using a strain range, $\Delta \varepsilon$, of ±0.5% and a temperature range, $\Delta T$, of 300°C with a minimum temperature, $T_{\text{min}}$, of 300°C and a maximum temperature, $T_{\text{max}}$, of 600°C. A saw-tooth waveform was used as shown in Figure 1a and the results for this test are shown in Figure 6a. The OP TMF test was carried out using a strain range, $\Delta \varepsilon$, of ±0.6% and the same temperature wave as the IP TMF test, the saw-tooth waveform was is shown in Figure 1b and the results for this test are shown in Figure 6b.
3. Definition of the material behaviour model

The Chaboche unified viscoplasticity model has been chosen to represent the uniaxial cyclic material behaviour of 316 stainless steel. The uniaxial form of the model is as follows:

\[ \dot{\varepsilon}_p = \left \{ \begin{array}{ll} \dot{f}, & x > 0 \\ 0, & x = 0 \\ -1, & x < 0 \end{array} \right \} sgn(\sigma - \chi) \]  
(1)

where, 
\[ sgn(x) = \left \{ \begin{array}{ll} 1, & x > 0 \\ 0, & x = 0 \\ -1, & x < 0 \end{array} \right \} \]
and
\[ f = |\sigma - \chi| - R - k \]  
(2)

where the elastic domain is defined by \( f \leq 0 \) and the inelastic domain by \( f > 0 \)

\[ \dot{\chi}_i = C_i \{ a_i \dot{\varepsilon}_p - \chi_i \dot{p} \} \]  
(3)

where \( i = 1, 2 \)

\[ \chi = \chi_1 + \chi_2 \]  
(4)

\[ \dot{R} = b(Q - R)\dot{p} \]  
(5)

\[ \sigma_v = Z\dot{p}^{\nu} \]  
(6)

\[ \dot{p} = |\dot{\varepsilon}_p| \]  
(7)

\[ \sigma = \chi + (R + k + \sigma_v)sgn(\sigma - \chi) = E(\varepsilon - \varepsilon_v) \]  
(8)

Equation (1), the viscoplastic flow rule, is the governing equation within the model. As can be seen from equations (2) to (8), all of the other model variables, such as those used for calculating both types of hardening (isotropic, \( R \) and kinematic, \( \chi \)) and viscous stress, \( \sigma_v \), are dependent on the value of \( f \).
plastic strain, $\varepsilon_p$, calculated from this viscoplastic flow rule. Equation (2) is the yield criterion for the model.

As previously mentioned, this model takes account of both kinematic hardening and isotropic hardening. Figure 7 shows the physical meaning of both types of hardening and the effect they have on the yield surface, both types of hardening are shown in three-dimensional (principle) stress space. When the stress state within the material causes the edge of the yield surface to be reached, kinematic hardening, implemented by equations (3) and (4), is represented as the movement of the yield surface, as shown in Figure 7a. Isotropic hardening, implemented by equation (5) represents the growth of the yield surface, as shown in Figure 7b.

![Figure 7. Schematic representations of hardening behaviour (a) Kinematic (b) Isotropic.](image)

Equation (6) defines the viscous stress, which takes the form of the widely used Norton creep law. Equation (7) shows that the variable, $p$, used in the calculation of many of the other variables is the accumulated plastic strain, $\varepsilon_p$.

The above model has been implemented in Matlab which is a top level programming language.

4. Identification of the material constants

In total, the material model requires the identification of 10 material constants. Within this section are brief descriptions of the methodologies used in calculating these constants, following the developments detailed in [11].

4.1. Initial yield stress, $k$ and Young’s modulus, $E$

From the initial experimental tensile curve, Young’s modulus, $E$ is taken as the gradient of the initial linear region. The initial yield stress, $k$, can be estimated as the stress value at the point at which the data begins to deviate from this initial linear region.

4.2. Isotropic hardening parameters, $Q$ and $b$

Equation (5) for the rate of isotropic hardening can be integrated with respect to time to give the following equation:

$$ R = Q \left(1 - e^{-b \varepsilon_p} \right) $$  \hspace{1cm} (9)

Equation (9) shows that as the accumulated plastic strain, $\varepsilon_p$, increases, $R$ exponentially approaches saturation to a value of $Q$. Therefore, assuming that the material hardening is entirely due to isotropic hardening and plotting $R$ against the accumulated plastic strain, the saturated value of $R$ is identified as $Q$, as shown in Figure 8. Then choosing a point roughly half way into the transient region of the hardening behaviour, shown circled in Figure 8, the corresponding values of $R$ and $p$ are identified. These values (along with the calculated value of $Q$) are then put into the following equation, which is simply a rearranged version of equation (9) for $b$. By choosing this point roughly half way into the transient region of the hardening behaviour, the value of $b$ calculated forces the model to go through
this point and with the model saturating at the value of $Q$, the result is a close model fit to the experimental data as shown in Figure 8.

\[
b = \left[ \ln\left(1 - \frac{R}{Q}\right) \right] \frac{1}{p}
\]  

(10)

Figure 8. Isotropic hardening variable $R$ versus $p$ for 316 stainless steel at 600°C.

4.3. Kinematic hardening parameters, $a_1$, $C_1$, $a_2$ and $C_2$

Equation (3) for the rate of kinematic hardening can be integrated, with respect to time, to give the following equations:

\[
\chi_1 = a_1 \left(1 - e^{-C_1 \epsilon_p}\right)
\]  

(11)

\[
\chi_2 = a_2 \left(1 - e^{-C_2 \epsilon_p}\right)
\]  

(12)

Substituting equations (11) and (12) into equation (8) gives:

\[
\sigma = a_1 \left(1 - e^{-C_1 \epsilon_p}\right) + a_2 \left(1 - e^{-C_2 \epsilon_p}\right) + R + k + \sigma_v
\]  

(13)

If the later stages of hardening are considered, it can be assumed that $\chi_1$ (and therefore $a_1$ and $C_1$) has a negligible effect on the hardening and therefore the kinematic hardening is dominated by $\chi_2$ ($a_2$ and $C_2$). Therefore, equation (13) can be simplified to:

\[
\sigma = a_2 \left(1 - e^{-C_2 \epsilon_p}\right) + R + k + \sigma_v
\]  

(14)

Differentiating equation (14) with respect to $\epsilon_p$, re-arranging and taking natural logs of both sides gives the following equation (assuming yield stress, $k$, and viscous stress, $\sigma_v$, to be constants):

\[
\ln\left(\frac{\partial \sigma}{\partial \epsilon_p} - \frac{\partial R}{\partial \epsilon_p}\right) = -C_2 \epsilon_p + \ln(a_2 C_2)
\]  

(15)

Therefore plotting $\ln\left(\frac{\partial \sigma}{\partial \epsilon_p} - \frac{\partial R}{\partial \epsilon_p}\right)$ vs. $\epsilon_p$ as shown in Figure 9 allows the identification of $C_2$ from the gradient and $a_2$ from the y-axis intercept. Similarly, $a_1$ and $C_1$ can be found, for the lower strain region, from equation (13), having already identified $a_2$ and $C_2$ [11].
Figure 9. Plot used in the calculation of the kinematic hardening material 
constants \( a_2 \) and \( C_2 \) for 316 stainless steel at 600°C.

In order to perform this fit to the data, it is necessary to obtain expressions for \( \frac{\partial \sigma}{\partial \varepsilon_p} \) and \( \frac{\partial R}{\partial \varepsilon_p} \), as 
functions of \( \varepsilon_p \), to use in equation (15). For the initial tensile curve (the first quarter cycle), \( \varepsilon = \varepsilon_p \), 
which can be substituted into equation (9). This expression can then be differentiated with respect to \( \varepsilon_p \), to give the following:

\[
\frac{\partial R}{\partial \varepsilon_p} = b Q e^{b \varepsilon_p} \tag{16}
\]

The calculation of \( \frac{\partial \sigma}{\partial \varepsilon_p} \) is more complicated. Taking \( \frac{d \sigma}{d \varepsilon_p} \), multiplying by \( \frac{dt}{dt} \) and \( \frac{d \varepsilon_r}{d \varepsilon_p} \), then 
rearranging gives the following:

\[
\frac{d \sigma}{d \varepsilon_p} = \frac{d \sigma}{d \varepsilon_r} \frac{1}{\dot{\varepsilon}_r} \dot{\varepsilon}_r \tag{17}
\]

Hence, expressions for \( \frac{d \sigma}{d \varepsilon_r} \) and \( \dot{\varepsilon}_r \) are required. The value of \( \dot{\varepsilon}_r \) is controlled during testing as 
hence is known. To obtain \( \dot{\varepsilon}_r \), Hooke’s law, \( \varepsilon = \frac{\sigma}{E} \), is substituted into the following equation for 
total strain:

\[
\varepsilon_r = \varepsilon_c + \varepsilon_p \tag{18}
\]

Rearranging equation (18) and differentiating it with respect to time gives the following:

\[
\frac{d \varepsilon_p}{dt} = \frac{d \varepsilon_r}{dt} - \frac{d \sigma}{dt} \frac{1}{E} \tag{19}
\]

Multiplying the final term in equation (19) by \( \frac{d \varepsilon_r}{d \varepsilon_r} \) and rearranging gives the following:
\[ \dot{e}_p = \dot{e}_T \left( 1 - \frac{1}{E} \cdot \frac{d\sigma}{d\varepsilon_T} \right) \]  

(20)

In order to obtain an expression for \( \frac{d\sigma}{d\varepsilon_T} \), a smoothing function is needed to eliminate complications caused by scatter in the experimental data, which could cause negative values of \( \frac{d\sigma}{d\varepsilon_T} \) to be obtained at some strain values. The smoothing function used in this case is the Ramberg-Osgood equation, i.e.:

\[ \varepsilon_T = \frac{\sigma}{\sigma_0} + \left( \frac{\sigma}{\sigma_0} \right)^{n_0} \]  

(21)

\[ \varepsilon_0 = \frac{\sigma_0}{E} \]  

(22)

Equation (22) can be substituted into equation (21) to give:

\[ \frac{E\varepsilon_T}{\sigma_0} = \frac{\sigma}{\sigma_0} + \left( \frac{\sigma}{\sigma_0} \right)^{n_0} \]  

(23)

which can then be differentiated with respect to \( \varepsilon_T \) to give:

\[ \frac{d\sigma}{d\varepsilon_T} = \frac{\sigma_0}{\varepsilon_0} \left( 1 + n_0 \left( \frac{\sigma}{\sigma_0} \right)^{n_0-1} \right) \]  

(24)

The Ramberg-Osgood constants, namely \( \varepsilon_0 \), \( \sigma_0 \) and \( n_0 \), can be found by rearranging and taking logs of both sides of equation (23) to give the following:

\[ \log(E\varepsilon_T - \sigma) = n_0 \log \sigma + (1 - n_0) \log \sigma_0 \]  

(25)

Therefore, plotting \( \log(E\varepsilon_T - \sigma) \) vs. \( \log \sigma \) allows the identification of \( n \) (gradient) and \( \sigma_0 \) (from the y-axis intercept). An example of this plot for a temperature of 600ºC is shown by Figure 10. Equation (22) can then be used to determine \( \varepsilon_0 \). Table 2 shows the Ramberg-Osgood constants calculated for the four temperatures.

![Figure 10](image_url)

**Figure 10.** Plot of \( \log(E\varepsilon_T - \sigma) \) vs. \( \log \sigma \) used in determining the constants
used in the Ramberg-Osgood equation.

Table 2. Ramberg-Osgood constants for 316 stainless steel at multiple temperatures.

<table>
<thead>
<tr>
<th>T (°C)</th>
<th>ε₀ (%)</th>
<th>σ₀ (MPa)</th>
<th>n₀</th>
</tr>
</thead>
<tbody>
<tr>
<td>300</td>
<td>0.21803</td>
<td>337.61</td>
<td>7.2409</td>
</tr>
<tr>
<td>500</td>
<td>0.20277</td>
<td>295.11</td>
<td>6.1627</td>
</tr>
<tr>
<td>550</td>
<td>0.21779</td>
<td>307.65</td>
<td>7.2846</td>
</tr>
<tr>
<td>600</td>
<td>0.19212</td>
<td>267.26</td>
<td>7.0468</td>
</tr>
</tbody>
</table>

4.4. Creep constants, Z and n

Typical values of Z and n have been taken from literature, such as Ryu [17] and Hyde [18-20]. The Matlab computer program for the model was run varying these constants around the typical values in order to obtain good fits to the model; this resulted in the constants presented in Table 2.

4.5. Material constants

Table 2 summarises the material constants identified from the isothermal test data for the four temperatures.

Table 3. Material constants for 316 stainless steel at multiple temperatures.

<table>
<thead>
<tr>
<th>T (°C)</th>
<th>k (MPa)</th>
<th>E (GPa)</th>
<th>b</th>
<th>Q (MPa)</th>
<th>a₁ (MPa)</th>
<th>C₁</th>
<th>a₂ (MPa)</th>
<th>C₂</th>
<th>Z (MPa.s^[1/n])</th>
<th>n</th>
</tr>
</thead>
<tbody>
<tr>
<td>300</td>
<td>39</td>
<td>154.84</td>
<td>39.46</td>
<td>32.76</td>
<td>119.1</td>
<td>5964.1</td>
<td>108.4</td>
<td>1001.6</td>
<td>179</td>
<td>10</td>
</tr>
<tr>
<td>500</td>
<td>32.5</td>
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<td>6472.6</td>
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<tr>
<td>550</td>
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<td>10</td>
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<tr>
<td>600</td>
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<td>27.43</td>
<td>80.06</td>
<td>7111.9</td>
<td>116</td>
<td>928.7</td>
<td>170</td>
<td>10</td>
</tr>
</tbody>
</table>

5. Model predictions

5.1. Comparison of model predictions to isothermal experimental results

Figure 11 to Figure 14 show comparisons of the experimental and model results for the isothermal cases of 550°C and 600°C for a strain range of ±0.3%. The figures show the monotonic tensile curve and first cycle, saturated cycle and hardening behaviour of the material. Figure 15 shows the complete predicted history (fifty loops) from the model for a temperature of 600°C, illustrating the predicted cyclic hardening phenomenon.

Figure 11. Comparison of model predictions of cyclic σ-ε behaviour to isothermal experimental data at 550°C. (a) Monotonic tensile curve and 1st loop (b) Saturated loop
Figure 12. Comparison of model predictions of hardening behaviour to isothermal experimental data at 550°C.

Figure 13. Comparison of model predictions of cyclic $\sigma$-$\varepsilon$ behaviour to isothermal experimental data at 600°C. (a) Monotonic tensile curve and 1st loop (b) Saturated loop

Figure 14. Comparison of model predictions of hardening behaviour to isothermal experimental data at 600°C.
Figure 15. Full $\sigma$-$\varepsilon$ model predictions for 50 loops under isothermal conditions at 600°C. (a) Full loops (b) Zoomed for maximum strain hardening behaviour

5.2. Comparison of model predictions to anisothermal experimental results

Table 3 shows that the material constants used in the Chaboche unified viscoplasticity model are generally temperature dependent. Therefore, one way of implementing the anisothermal condition within the model is to identify expressions for the constants as functions of temperature and inputting these expressions to the model rather than single values. This allows each of the constants to be calculated at the beginning of each time increment depending on the value of temperature at the beginning of that time increment. These values are then used in the calculation of the subsequent values of plastic strain, stress, etc, hence creating a temperature-dependent (TMF) material model.

Figure 16 and Figure 17 show the experimental/model comparisons for the two TMF test conditions.

Figure 16. Comparison of model predictions of cyclic $\sigma$-$\varepsilon$ behaviour to in-phase TMF experimental data at high strain range. (a) Monotonic tensile curve and 1st loop (b) Saturated loop
6. Conclusion

A unified material model, which includes both (non-linear) isotropic and kinematic hardening behaviour as well as viscoplasticity phenomena, such as rate-dependency, has been successfully implemented in Matlab. A programme of isothermal and anisothermal thermo-mechanical tests have been conducted using induction coil heating, with the temperature uniformity controlled to within ±10°C up to temperatures of 600°C. The isothermal tests have been employed to identify the material constants for the material model at different temperatures between 300°C and 600°C. The experimental data has also been employed to validate the Matlab implementation of the unified viscoplasticity model, showing excellent model to test correlation for the isothermal tests considered. Interpolation of the temperature-dependency of the material constants, for application to the anisothermal simulations has also shown reasonably good correlation, although the model appears to slightly under-predict the peak tensile stresses during forward plastic deformation and over-predict the intermediate to high compressive stresses during reverse plastic deformation. It is also worth noting that the material constants were obtained only from the ±0.3% strain range (isothermal) data, due to the use of the stepped strain-range technique to minimise specimen usage, whereas the TMF experimental data was obtained at strain ranges of ±0.5% and ±0.6%, the model predictions therefore represent an extrapolation in terms of the strain range used. Further testing to characterise TMF behaviour of the material at the strain range at which the material constants were obtained is expected to improve the anisothermal predictions.

A parallel project is concerned with development of optimisation techniques for automating and refining the process of material constant identification.

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