# Microstructural Evolution and Constitutive Equations of Inconel 718 Alloy under Quasi-static and Quasi-dynamic Conditions

M. Azarbarmas<sup>a,b\*</sup>, M. Aghaie-Khafri<sup>a1</sup>, J.M. Cabrera<sup>b</sup>, J. Calvo<sup>b</sup>

<sup>a</sup> Faculty of Materials Science and Engineering, K.N. Toosi University of Technology, Postal Code: 1999143344, Tehran, Iran

<sup>b</sup> Departament de Ciència dels Materials i Enginyeria Metal·lúrgica, ETSEIB – Universitat Politècnica de Catalunya, Av. Diagonal 647, 08028 Barcelona, Spain

## Abstract

The deformation behavior of IN718 superalloy was investigated using the hot compression tests in the temperature range of 950-1100  $^{\circ}$ C, and strain rates covering the quasi-static to the quasidynamic regions (0.001-10 s<sup>-1</sup>). The shape of flow curves as well as the corresponding work hardening rates analysis was utilized to reveal the dynamic recrystallization (DRX) phenomena. DRX was the dominant restoration mechanism in the whole temperature and strain rate domains, which was characterized by the optical and EBSD images. Extended flow softening was observed at high strain rates due to the adiabatic heating and dislocations interaction. In addition to the assessment the capability of Sellars equations, a new constitutive equation based on the multiple variable regression analysis was proposed for modeling the peak stress as a function of

<sup>\*</sup> Corresponding author.

E-mail addresses: mazarbarmas@ut.ac.ir (M. Azarbarmas)

strain rate and temperature. Besides the simple form of the proposed model, it has a good accuracy for predicting the peak stress.

**Keywords:** Inconel 718; Hot compression test; Constitutive equation; Dynamic recrystallization; EBSD.

## 1. Introduction

The Nickel-based superalloy IN718 is a precipitation hardenable alloy with high corrosion resistance [1,2], excellent resistance to oxidation [3], high strength [4], adequate ductility and toughness [5], and good machining performance [6]. IN718 is an important material used for modern aero-engines, gas-turbines, extrusion dies, etc. [7–15] Control of the thermomechanical processing is crucial for obtaining favorable microstructure and properties. The hot deformation behavior of IN718 has been widely studied over the past decade using hot compression tests [6,16–19]. Considering Sellars and Tegart equations, some models have been proposed in the form of an hyperbolic-sine [20–22], an exponential [23] and a power law [16,24] for hot compression of IN718. Furthermore, J.M. Zhang et al. [25] have formulated experimental data by considering the shape of the flow stress vs. deformation parameters curves to develop a mathematical model. Recently, M.Z. Hussain et al. [26] have proposed a constitutive relationship for thermo-mechanical processing of IN718 through double multivariate nonlinear regression. Considering hot deformation of IN718, it is worth noting that it is essential to develop comprehensive equations for a wide range of temperature and strain rate.

IN718 is a material with relatively low stacking fault energy [22], in which the presence of solute atoms and phases reduce the mobility of dislocations. Therefore, the main restoration process is dynamic recrystallization (DRX), although the dynamic recovery (DRV) softening plays also a

competitive role. DRX is an important mechanism that controls the microstructure evolution of the IN718 during the hot working of the material, and indeed, there are numerous works in the literature which have explored the parameters affecting DRX in IN718 [17,27–32]. Due to the enormous influence of DRX on the microstructure and properties after processing, the evaluation of DRX in terms of deformation parameters is vital.

The main goal of the present work is to investigate the hot working behavior of IN718 during the hot compression test at quasi-static and quasi-dynamic conditions. Besides, general constitutive equations and a new model based on the multivariable regression were employed to obtain the hot working constants of the alloy and assess which of them are most appropriate in all conditions.

# 2. Experimental procedure

## 2.1. Material

The starting material in this study was wrought IN718 alloy. Chemical composition of the material is shown in Table 1. The specimens were solution treated at 1025 for 1 h, followed by cold-water quenching. The initial microstructure comprises equiaxed grains and annealing twins, as is shown in Fig. 1. The average grain size (without twins) was measured to be about 45  $\mu$ m by the linear interce °C pt method.

## **2.2.** Compression test

Cylindrical specimens with a diameter of 5 mm and a height of 10 mm were used for hot compression tests. Baehr DIL-805 deformation dilatometer was used for isothermal compression

tests. Compression tests were conducted up to true strain of about 0.7 under different temperatures (950, 1000, 1050 and 1100 °C) and various strain rates (0.001, 0.01, 0.1, 1 and 10 s<sup>-1</sup>), for covering the quasi-static and quasi-dynamic condition. A K-type thermocouple was spotwelded at the middle of the specimen to accurately control the heating and cooling rates and measure the specimen temperature. Also, argon gas was utilized as the shielding and quenching gas in the machine. Hot compression tests were conducted in an argon atmosphere. In order to minimize the frictions during the hot deformation and to avoid the adhesion, a Molybdenum foil with a thickness of 0.1 mm was used between anvils and specimen surface.

#### 2.3. Microstructural studies

Hot deformed samples were sectioned parallel to the compression direction. Samples for optical metallographic examination were prepared by mechanically polishing and etching with a solution consisting of 50 ml HCl, 50 ml ethanol and 7.5 g CuCl<sub>2</sub>. For EBSD *analysis, specimens* were ground with SiC papers, mechanically polished with diamond slurries and then were polished with 0.04  $\mu$ m colloidal silica solution for one hour. A Zeiss Ultra Plus analytical field emission gun scanning electron microscope (FEG-SEM) equipped with an EBSD detector, provided by HKL Technology, was used for EBSD studies with a step size of 0.25  $\mu$ m. analyzing and displaying the EBSD data were done by using the Channel5 software package.

# 2.4. Correction of flow curves

It is necessary to use correction factors for eliminating the effect of interfacial friction. A simplified analysis of the barrel compression test based on the upper-bound theory [33] was used to remove the friction effects on the flow curves.

Generally, the occurrence of DRX can be identified from the appearance of a peak stress in the flow curves. Nevertheless, in some cases detection of this peak is difficult under hot working conditions. The initiation of DRX can also be identified from inflection point of the strain hardening rate against stress curves [34,35]. The first derivative of the true stress versus true strain yields the work hardening rate ( $\theta$ ). The  $\theta$  values were obtained by averaging the slopes of two adjacent points for each data point using Origin program as follows:

$$\frac{1}{2} \left( \frac{y_{i+1} - y_i}{x_{i+1} - x_i} + \frac{y_i - y_{i-1}}{x_i - x_{i-1}} \right)$$
(1)

where x and y are the coordinates of each data point. For solving noise problems, a curve smoothing by fitting a seven to nine-order polynomial to the stress-strain curves was used to eliminate the fluctuations during differentiation[36,37]. In this study, following the removal of the elastic portions of curves, each curve was friction-corrected and then was smoothed with a high-order polynomial.

# 3. Results and discussion

#### 3.1. Corrected flow curves

The friction-corrected flow curves obtained for different temperatures and strain rates are presented in Fig. 2. Most of samples exhibit typical DRX flow curves with a clear single peak stress followed by a gradual decrease towards a steady state stress. Having a single peak behavior is the main characteristic of discontinuous dynamic recrystallization[35], and shows that recrystallization initiates in new cycles before completion of the current cycles [36]. Thus, different grains are at different state of recrystallization development during deformation and the single peak flow curve is the averaged flow stress of grains at various stages of recrystallization.

As can be observed in Fig. 2, the peaks of flow curves become less obvious with decreasing strain rate and increasing temperature. The abnormal stress drop beyond the peak stress in the case of samples deformed at  $10 \text{ s}^{-1}$  strain rate is due to the adiabatic deformation heating. When hot compression is performed at strain rates of 10 s<sup>-1</sup> and higher, the degree of temperature rise due to adiabatic heating is high and the heat cannot dissipate to the surrounding environment during the short time periods of the tests [38]. Additionally, considering that the deformation process is very fast, it is almost impossible for the testing machine to compensate the temperature deviation during the compression test. On DIL-805 deformation dilatometer, the instantaneous temperature in the stable center region of specimen was measured by a welded thermocouple during hot deformation. Fig.3a shows the adiabatic deformation heating for the specimen deformed at high strain rate of 10 s<sup>-1</sup> and 1050 °C. Similar behavior at 10 s<sup>-1</sup> strain rate has been reported by H. Mirzadeh et al. for hot compression of steels [36,39]. It can be observed that the increase in temperature following the peak point is significantly high owing to the adiabatic heating. The temperature rise results in reduction in the flow stress of the sample, i.e., the peak stress (or the stress softening) in this condition is related to the temperature increment rather than to the onset of DRX. Analogous plots at T = 1050  $^{\circ}$ C and 10 s<sup>-1</sup> strain rate are presented in Fig.3b. This figure shows that the adiabatic heating during deformation is negligible for samples compressed at strain rates lower than  $10 \text{ s}^{-1}$ .

It is clear from Fig. 2 that the plastic region of the flow curve consists of three stages: i) work hardening, ii) softening and iii) steady state. The first stage is the work hardening region, in which dislocation density continuously increases and the flow stress quickly increases with the increase of the strain. In the softening region, the dislocation density reaches a critical level, and DRX occurs. During the nucleation and growth of recrystallized grains the number of

dislocations is reduced and thereby the flow stress decreases. Finally, at the steady state, there is a balance between work hardening and dynamic softening.

#### 3.2. Microstructural characteristics

Hot deformation at different temperatures and strain rates results in grain elongation, boundaries serration, substructure formation and finally the onset of DRX after a critical strain. As an illustrative example, the effects of deformation temperature on the microstructure of the specimens under the strain rate of 0.01 s<sup>-1</sup> is shown in Fig. 4. It can be observed that for a given nominal strain of 0.7, the fraction of recrystallized grains increases with the increase of the deformation temperature. Considering that the occurrence of DRX, at a given initial grain size, is dependent on the distribution and of dislocation density [21], and knowing that the increasing of the deformation temperature decreases the critical dislocation density, the critical strain for the onset of DRX decreases. As depicted in Fig. 4, with increasing the temperature the microstructures transform from a duplex or "necklace" to a completely recrystallized structure and the recrystallized grain size increases. The recrystallized grain growth is due to high mobility of the grain boundaries at high temperatures.

Fig. 5 depicts the influences of strain rate on the microstructures of deformed specimens at the deformation temperature of 1050 °C. It is clear that the recrystallized grain size decreases with increasing strain rate. Moreover, the fraction of DRX decreases with increasing the strain rate. The exceptional case for the relationship of grain size and the fraction of DRX with strain rate is the specimen compressed at strain rate of 10 s<sup>-1</sup>, which shows higher grain size and greater fraction of DRX in comparison with the specimen deformed at strain rate of 1 s<sup>-1</sup>. The promotion of DRX at high strain rate of 10 s<sup>-1</sup> can be ascribed to the adiabatic heating and increased rate of

dislocation accumulation. Adiabatic heating within the specimen facilitates DRX nucleation and grain boundary migration and thereby stimulates DRX process. On the other hand, due to the strong dislocation–dislocation interaction and weak recovery during plastic deformation at high strain rate, the dislocation density increases. *Distributions of sub-boundaries inside the elongated grains have been compared for samples deformed at temperature of 1100* °C and strain rates of 10 *and 1* s<sup>-1</sup> *in Fig. 6. The* more density of *low-angle grain boundaries for sample compressed at strain rate of 10* s<sup>-1</sup> *approves the increasing dislocation density with increasing the strain rate.* The increased dislocation density decreases the recrystallization temperature noticeably [40]. Therefore, the high volume fraction and grain size of recrystallized grains at strain rate of 10 s<sup>-1</sup> are resonable.

In a single-peak DRX process, nucleation occurs essentially along existing grain boundaries (necklace mechanism). The growth of new grains is *limited* by the concurrent deformation as a result of decreased driving force for their subsequent growth due to rising the dislocation density [41]. The DRX process continues until the completion of the first layer of necklace to cover the entire grain boundary. Afterward, the subsequent layers form at the recrystallization front between the recrystallized and un-recrystallized portions.

The OIM band contrast microstructures of annealed sample and compressed at  $1000^{\circ}$ C-0.1 s<sup>-1</sup> are *presented* in Fig. 7. It is evident from Fig. 7a that the initial annealed microstructure (preceded to hot deformation) consisted of equiaxed grains with a large quantity of annealing twin boundaries and negligible fraction of low angle boundaries. The equiaxed grains are the result of static recrystallization and grain growth during the annealing heat treatment. There are two types of twin boundaries in the structure, i) the straight and ii) the curved twin boundaries which are *indicated with white and yellow arrows, respectively, shown in Fig.7a*. It is believed that the

straight twin boundaries are coherent and the curved are incoherent [42]. As expected for a material with low stacking fault energy [43] the presence of this high fraction of twin boundaries in the microstructure shows that formation of annealing twins is an important phenomenon during the recrystallization of this alloy. Twin domains are favorable nucleation sites for DRX due to their high stored deformation energy [44].

Fig. 7b *depicts plenty of* sub-boundaries in the deformed and un-recrystallized matrix of the sample deformed at  $1000^{\circ}$ C-0.1 s<sup>-1</sup>, which is a result of dislocation generation and dislocation boundary formation during the plastic deformation and dynamic recovery. However, the new DRX grains consume the deformed microstructure and thereby decrease the fraction of low angle boundaries. Besides, most of the twin boundaries in the matrix have been diminished. Some of the pre-existing twins have lost their twin characteristics and curved due to the effects of concurrent deformation [43,45], showed by yellow arrow in Fig. 7b. *It is noteworthy that* a large fraction of newly generated boundaries after the initiation of DRX have twin relationship. It is obvious *from Fig* 7b that the dislocation sub-boundaries behind the bulged parts are free of twins. Therefore, it can be pointed out twins inside DRX grains form during the growth of DRX grains. This is in line with the other reported *observations* [46].

The typical macrograph and micrograph of specimens deformed under 1050 °C-10 s<sup>-1</sup> are shown in Fig. 8a and b, respectively. These images show large amount of local deformation bands forming about 45 ° angles to the compression direction and indicate the existence of the instability in the form of unstable flow due to the flow localization at high strain rates. The same phenomenon has also been found in compression tests of superaustenitic stainless steel and has been interpreted as a result of flow localization [47].

#### **3.3.** Constitutive equations based on Sellars models

Several empirical equations have been presented to characterize the hot deformation behavior and to obtain constitutive equations of alloys. These equations are commonly employed to define *the flow stress as a function of strain rate and deformation temperature*[48,49]. Generally, a constitutive equation is determined by formulating experimental data [25].

Hot deformation can be conceived as a thermally-activated process which is modeled with strain rate equations like to those used in creep modeling [50]. The combined effect of temperature and strain rate is classically unified through the Zener-Hollomon parameter (*Z*), whose physical interpretation is the strain rate factor compensated by temperature.

$$Z = \operatorname{seep}\left(\frac{Q}{RT}\right) \tag{2}$$

where & is strain rate (s<sup>-1</sup>), Q is hot deformation activation energy (kJ/mol), R is the gas constant (8.314 kJ/mol K<sup>-1</sup>), and T is deformation temperature (K).

Concerning developing constitutive equations, the Z parameter can be associated with the flow stress in different ways. Three types of equations are mostly used according to the deformation conditions, involving exponential, power and hyperbolic sine functions [51–53].

$$Z = \mathscr{E} \exp\left(\frac{Q}{RT}\right) = f(\sigma) = \begin{cases} A_1 \exp(\beta\sigma) \\ A_2 \sigma^{n'} \\ A_3 [\sinh(\alpha\sigma)]^n \end{cases}$$
(3)

where,  $A_1$ ,  $A_2$ ,  $A_3$ , n', n,  $\beta$  and  $\alpha$  ( $\approx \beta/n'$ ) are experimentally determined temperatureindependent material constants. In these equations, the flow stress is related to the strain rate and temperature of deformation. Nevertheless, such flow stress definition must be considered as rate equations rather than constitutive equations because there is no dependence on strain in the equations. Typically the latter equations can be applied to characteristic stresses such as steady state or peak stress. The method used for obtaining the characteristic stresses, peak and steady state stress, is illustrated in Fig. 9. It appears from Fig. 10 that the peak stress and the steady-state stress have linear relationship especially for samples compressed at low strain rates. Although in the in literature, there are reports of using both the peak stress [54] and the steady-state stress [16] for formulating the constitutive equations, one should notice that it is not often possible to determine the steady state stress accurately and the adiabatic heating can drop the flow stress at high strain rates. Therefore, it seems that it is better to use the peak stress in constitutive equations rather than steady state stresses.

Eq. (2) can be rearranged to yield the following equations:

$$\ln \mathscr{E}_{+}\left(\frac{Q}{R}\right)\frac{1}{T} = \begin{cases} \ln A_{1} + \beta\sigma\\ \ln A_{2} + n'\ln\sigma\\ \ln A_{3} + n\ln\left[\sinh\left(\alpha\sigma\right)\right] \end{cases}$$
(4)

 $\beta$ , n' and n can also be derived by partial differentiation of Eq. (3) at a given deformation temperature:

$$\begin{cases} \beta = \left[\frac{\partial \ln \mathscr{X}}{\partial \sigma}\right]_{T} \\ n' = \left[\frac{\partial \ln \mathscr{X}}{\partial \ln \sigma}\right]_{T} \\ n = \left[\frac{\partial \ln \mathscr{X}}{\partial \ln \left[\sinh(\alpha\sigma)\right]}\right]_{T} \end{cases}$$
(5)

The slopes of the curves of  $\ln \mathscr{E}$  versus  $\sigma_p$ ,  $\ln \sigma_P$  and  $\ln[\sinh(\alpha\sigma)]$  can be employed for determining the values of  $\beta$ , n' and n, respectively. The stress multiplier  $\alpha$  can be estimated by knowing the values of  $\beta$  and n' ( $\alpha \approx \beta / n'$ ) [55], which is obtained to be 0.006. Aforementioned

plots are presented in Fig. 11. The average values of  $\beta$ , n' and n were calculated by linear regression of the corresponding curves and are listed in Table 2. The adjusted R<sup>2</sup> error values presented in Table 2 reveal that the linear regression results are acceptable.

Partial differentiation of Eq. (4) at constant strain can be used for calculating the value of deformation activation energy (Q). The necessary expressions are as follows:

$$Q = \begin{cases} R\beta \left[ \frac{\partial \sigma}{\partial (1/T)} \right]_{\&} \\ Rn' \left[ \frac{\partial \ln \sigma}{\partial (1/T)} \right]_{\&} \\ Rn \left[ \frac{\partial \ln \left[ \sinh (\alpha \sigma) \right]}{\partial (1/T)} \right]_{\&} \end{cases} \end{cases}$$
(6)

Based on Eq.(6) the slope of the plots of  $\sigma_p$ , ln  $\sigma_p$  and ln sinh( $\alpha\sigma_p$ ) versus the reciprocal of the absolute temperature can be utilized for calculating the value of Q. The required plots are shown in Fig. 12. As is shown in Table 3, the average values of 442, 409, and 437 kJ/mol were obtained for activation energy based on the exponential, power and hyperbolic sine laws, respectively. It can be observed that all the values are close to each other. The average activation energy of hot deformation was considered to be 429 kJ/mol which is comparable to the values reported for hot deformation of IN718: 443 kJ/mol by Y. Wang et al. [21] and 429 kJ/mol by L. Cheng et al. [56]. However, the activation energy is significantly higher than the activation energy for self-diffusion of nickel (278 kJ/mol [57]). The higher Q value can be ascribed to the influences of alloying elements which pin the motion of the dislocations and grain boundaries. The final obtained equations after *substituting Q, n, ń, a and B values are as follow:* 

$$\ln \& + \frac{51600}{T} = \begin{cases} \ln A_1 + 0.026\sigma \\ \ln A_2 + 5.029 \ln \sigma \\ \ln A_3 + 3.462 \ln [\sinh(0.005\sigma)] \end{cases}$$

# 3.4. Peak Stress as a Function of the Zener–Hollomon Parameter

Considering Eq. (3), the curves of ln Z against  $\sigma_p$ , ln  $\sigma_p$ , or ln sinh( $\alpha\sigma_p$ ) were plotted, Fig. 13, and used to model the relationship between Z and  $\sigma_p$ . Using a linear regression, the following equations with new constants were derived:

(7)

$$Z = \sec \left(\frac{429152}{\text{RT}}\right) = \begin{cases} (3.16 \times 10^{13}) e^{(0.027\sigma_{\rm p})} \\ 61023\sigma_{\rm p}^{4.957} \\ (2.57 \times 10^{15}) \left[ \sinh(0.0054\sigma_{\rm p}) \right]^{3.562} \end{cases}$$
(8)

Considering Table 4, the power law equation has the highest adjusted  $R^2$  value. Besides, this equation is simple and easy to use. Therefore, it is preferred to use the power law for modeling the hot deformation behavior of IN718 in the temperature range of 950-1100 °C and strain rate range of 0.001-10 s<sup>-1</sup>. However, the hyperbolic sine law has also a good fit and the exponential equation is not precise in the present investigation. Finally, in the order to increase the accuracy of peak stress calculation in IN718 alloy under the deformation condition used in this study the simplified following equations can be used:

(9)  

$$\sigma_{\rm p} = 37.04 Ln Z - 1151$$

$$\sigma_{\rm p} = 185 \times Sinh^{-1} (12475 \times Z^{0.28})$$

$$\sigma_{\rm p} = 0.108 \times Z^{0.202}$$

#### 3.5. A new model based on multivariable regression

Based on the experimental data of hot compression tests, a non-linear constitutive model is constructed by multiple linear regression analysis. Fig. 11a shows that *a linear relationship exists between*  $Ln\sigma_p$  and  $Ln\mathscr{R}$  at constant temperature. On the other hand, it is clear from Fig. 12b that at a given strain rate there is a linear relation between  $Ln\sigma_p$  and (1/T). By simple linear regression analysis of Fig 11a and 12b, the following regression equations can be derived:

$$Ln\sigma_P = A Ln \mathscr{S} + \sigma_P(T) \tag{10}$$

$$Ln\sigma_{P} = B\left(\frac{1}{T}\right) + \sigma_{P}\left(\mathscr{B}\right) \tag{11}$$

It is important to note that the slope of  $Ln \sigma_P$  - Ln & curve is independent of temperature and that of  $Ln \sigma_P - \frac{1}{T}$  curve is independent of strain rate in Fig 11a and 12b, respectively. It means that A and B in Eq. (10) and (11) are constant at all temperatures and strain rates. Thus, the following expression can be proposed for the peak stress:

$$Ln\sigma_{P} = A Ln \mathscr{S} + B\left(\frac{1}{T}\right) + C \tag{12}$$

This equation is in the form of  $y = y_0 + ax_1 + bx_2$ . Therefore, the *multivariate linear regression* can be used for finding the A, B and C constants. Based on the constructing data, the peak stress model or the constitutive equation for the hot deformation of IN718 is obtained as follows:

$$Ln\sigma_{p} = 0.199 Ln\mathscr{A} + \frac{10709.6}{T} - 2.458 \tag{13}$$

Fig. 14a shows 3D illustration of the experimental and calculated peak stress based on the Eq. (13). For better assessing the prediction capabilities of the model, results of hot compression

tests of IN 718 reported by H. Y. Zhang et al.[22] and J. M. Zhang et al. [25]were utilized. The important criterion in selection of data from the literature was that the strain rate or deformation temperature of those works should differ from that of our work. The experimental peak stress of IN 718 obtained at deformation temperatures of 980[25] and 1060 °C[22] under various strain rates are presented in Fig. 14 b. This figure shows that the prediction of the model is very close to that of the test case, especially for higher temperatures. The adjusted  $R^2$  of 0.962 and 0.975 for deformation temperatures of 980 and 1060 °C, respectively, shows that the established constitutive relationship is capable of describing the peak stress of the alloy at different strain rates and deformation temperatures. By some algebraic manipulations of Eq. (12), the following equation is derived.

$$C'\sigma_P^{\frac{1}{A}} = \mathscr{E}\left(\frac{B}{AT}\right)$$
(14)

By comparing Eq. (14) with the second expression of Eq. (3), A and B constants can be physically described. It can be observed that 1/A and BR/A represent n' and activation energy (Q), respectively.

The advantage of the new proposed constitutive equation in comparison to the Sellars-Tegart based equations is that the propos equation is simple and can be easily obtained. Moreover, there is no need for calculating the activation energy of the hot deformation.

#### Conclusions

A step-by-step data analysis of the hot compression behavior of IN718 alloy at temperatures ranging from 950 to 1100  $^{\circ}$ C and strain rates of 0.001 to 10 s<sup>-1</sup> resulted in the following principal conclusions:

-The inflection analysis in the work hardening rate versus stress plots as well as optical and EBSD images confirmed that the flow curves exhibited typical DRX behavior.

-The specimen compressed at high strain rate showed an exceptional DRX behavior and extended softening because of the adiabatic heating and strong interaction between the dislocations.

- The microstructure of the alloy at high temperatures and high strain rates had a large amount of local deformation bands forming about  $45^{\circ}$  angles to the compression direction showing an instable behavior.

-Among Sellars *constitutive* equations, the power law constitutive equation has the highest correlation coefficient and also is simple and easy to use.

-A new multivariable regression based model was proposed for obtaining constitutive equation of the material. The simple proposed model showed a good accuracy for predicting the peak stress as a function of strain rate and temperature.

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# **Figures Captions**

Fig. 1. Optical micrograph of the studied IN718 before hot deformation.

Fig. 2. True stress-true strain curves obtained in hot compression of IN718 at different strain

rates and temperatures of (a) 950, (b) 1000, (c) 1050 and (d) 1100 °C.

Fig. 3. (a) True stress and temperature versus true strain curves for samples deformed at 1050 °C

and strain rate of (a) 10 and (b)  $1 \text{ s}^{-1}$ .

Fig. 4. Microstructure of IN718 alloy deformed at strain rate of 0.01 s<sup>-1</sup> and temperature of (a) 950, (b) 1000, (c) 1050 and (d) 1100 °C; showing DRX microstructures.

Fig. 5. Microstructure of IN718 alloy deformed at strain rate of (a) 0.001, (b) 0.01, (c) 0.1, (d) 1 and (e) 10 s<sup>-1</sup>, and 1050 °C; showing DRX microstructures.

Fig. 6. The OIM band contrast microstructures for samples deformed at  $1100^{\circ}$ C and strain rate of (a) 0.1 s<sup>-1</sup> and (b) 10 s<sup>-1</sup>. The high and low angle boundaries are shown as black and white lines, respectively.

Fig. 7. The OIM band contrast microstructures for (a) annealed sample and (b) sample deformed at  $1000^{\circ}$ C-0.1 s<sup>-1</sup>; the high and low angle boundaries are shown as black and white lines, respectively;  $\Sigma$ 3 twin boundaries are shown as red lines.

Fig. 8. (a) Macrograph and (b) micrograph of IN 718 alloy deformed at 1050 °C-10 s<sup>-1</sup>.

Fig. 9. (a) Some of work hardening rate-true stress curves in various conditions; (b) Method used for determination of peak and steady state stresses.

Fig. 10. The linear relationship between (a) the peak stress and steady state stress at various strain rates, and (b) variations of  $\ln \sigma$  and  $\ln \vartheta$ .

Fig. 11. Plots used for calculation of (a) n', (b)  $\beta$  and (c) n values.

Fig. 12. Plots of (a)  $\sigma_{p}$ , (b) ln  $\sigma_{p}$ , and (c) ln sinh( $\alpha\sigma_{p}$ ) versus the reciprocal of absolute temperature, used for obtaining the values of Q.

Fig. 13. Plots of ln Z against (a)  $\sigma_P$ , (b) ln  $\sigma_P$ , and (c) ln sinh( $\alpha\sigma_P$ ).

Fig. 14. Evaluation the proposed model; (a) red points are the experimental values obtained in this work; (b) triangular and square denote the experimental data taken from Ref. [25] and Ref. [22], respectively, and the solid lines the calculated values using the proposed equation.

# **Tables Captions**

Table 1. Chemical composition of IN 718 (wt.%)

Table 2. Calculated values required to find the average values of  $\beta$ , n' and n (Eq. (5)) from curves of ln & versus  $\sigma_p$ , ln  $\sigma_P$  and ln  $\lceil \sinh(\alpha \sigma) \rceil$ 

Table 3. Calculated values required to find the values of activation energy from plots of  $\sigma_P$ , ln  $\sigma_{\rm p}$  and ln sinh( $\alpha\sigma_{\rm p}$ ) versus 1/T

Table 4. Calculated values of  $\beta$ , n' and n from ln Z against  $\sigma_p$ , ln  $\sigma_p$ , or ln sinh( $\alpha\sigma_p$ ) curves

Table 1. Chemical composition of IN 718 (wt.%)

Ni	Cr	Fe	Nb+Ta	Мо	Ti	Al	Со	SI	Mn	Cu	С
54.00	18.21	17.29	5.35	2.98	0.91	0.66	0.24	0.08	0.07	0.07	0.03

Table 2. Calculated values required to find the average values of  $\beta$ ,  $\acute{n}$  and n (Eq. (5)) from f ln  $\dot{\varepsilon}$  versus  $\sigma_{\rm e}$  ln  $\sigma_{\rm P}$  and ln [sinh(ad

curves	of ln	Ė	versus	$\sigma_n$	ln	$\sigma_P$	and	ln	[sinh(	ασ	)]	
	0, 111	C	1015005	• <i>p</i> ,		Οr			[000000	(000	71	

T (0C)	Ex	ponential law		Power law	Hyperbolic sine law		
I (°C)	β	Adjusted R <sup>2</sup> of β	n'	Adjusted R <sup>2</sup> of n'	n	Adjusted R <sup>2</sup> of n	
950	0.020	0.9355	4.838	0.9948	2.987	0.9644	
1000	0.027	0.9838	5.501	0.9914	3.768	0.9990	
1050	0.025	0.9463	4.931	0.9926	3.389	0.9908	
1100	0.031	0.9257	4.846	0.9988	3.704	0.9905	
Average	0.026	0.9478	5.029	0.9944	3.462	0.9862	

*Table 3. Calculated values required to find the values of activation energy from plots of*  $\sigma_P$ *, ln*  $\sigma_P$ 

	Exp	onential la	W	I	Power law		Hyperbolic sine law		
Strain Rate (s <sup>-1</sup> )	$\begin{array}{lll} Slope \ of & Adjuste \\ \sigma_{P} - \frac{1}{T} & d \ R^{2} of \\ & slope \end{array}$		Q (kJ/mol)	Slope of $\ln \sigma_{\rm P} - \frac{1}{T}$	Adjusted R <sup>2</sup> of slope	Q (kJ/mol)	Slope of ln sinh $(\alpha \sigma_p) - \frac{1}{T}$	Adjusted R <sup>2</sup> of slope	Q (kJ/mol )
0.001	1006467	0.9604		11525	0.9859		12630	0.9855	
0.01	1474900	0.9878		10597	0.9941		13073	0.9980	
0.1	2484764	0.9689		10902	0.9972		17219	0.9864	
1	2190106	0.9936		7792	0.9999		14117	0.9966	
10	3064037	-		8106	-		18793	-	
Averag e	2044055	0.9777	442	9784	0.9943	409	15166	0.9916	437

and  $ln sinh(\alpha \sigma_P)$  versus 1/T

Table 4. Calculated values of  $\beta$ ,  $\acute{n}$  and n from  $\ln Z$  against  $\sigma_P$ ,  $\ln \sigma_P$ , or  $\ln \sinh(\alpha \sigma_P)$  curves

Expo	nential law	Р	ower law	Hyperbolic sine law		
β	Adjusted R <sup>2</sup> of β	n′	Adjusted <b>R</b> <sup>2</sup> of n'	n	Adjusted R <sup>2</sup> of n	
0.027	0.9423	4.957	0.9935	3.562	0.9889	



Fig. 1. Optical micrograph of the studied IN718 before hot deformation.



Fig. 2. True stress-true strain curves obtained in hot compression of IN718 at different strain rates and temperatures of (a) 950, (b) 1000, (c) 1050 and (d) 1100 °C.



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Fig. 5. Microstructure of IN718 alloy deformed at strain rate of (a) 0.001, (b) 0.01, (c) 0.1, (d) 1 and (e)  $10 \text{ s}^{-1}$ , and  $1050 \degree$ C; showing DRX microstructures.



Fig. 6. The OIM band contrast microstructures for samples deformed at  $1100^{\circ}$ C and strain rate of (a) 0.1 s<sup>-1</sup> and (b) 10 s<sup>-1</sup>. The high and low angle boundaries are shown as black and white lines, respectively.



Fig. 7. The OIM band contrast microstructures for (a) annealed sample and (b) sample deformed at 1000°C-0.1 s<sup>-1</sup>; the high and low angle boundaries are shown as black and white lines, respectively;  $\Sigma$ 3 twin boundaries are shown as red lines.



Fig. 8. (a) Macrograph and (b) micrograph of IN 718 alloy deformed at 1050 °C-10 s<sup>-1</sup>.



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