On the Influence of Nb/Ti Ratio on Environmentally-Assisted Crack Growth in High-Strength Nickel-Based Superalloys

A.A.N. NÉMETH, D.J. CRUDDEN, D.M. COLLINS, V. KUKSENKO, C.H. LIEBSCHER, D.E.J. ARMSTRONG, A.J. WILKINSON, and R.C. REED

The effect of Nb/Ti ratio on environmentally-assisted crack growth of three prototype Ni-based superalloys is studied. For these alloys, the yield strength is unaltered with increasing Nb/Ti ratio due to an increase in grain size. This situation has allowed the rationalization of the factors influencing damage tolerance at 700 °C. Primary intergranular cracks have been investigated using energy-dispersive X-ray spectroscopy in a scanning transmission electron microscope and the analysis of electron back-scatter diffraction patterns. Any possible detrimental effect of Nb on the observed crack tip damage due to Nb-rich oxide formation is not observed. Instead, evidence is presented to indicate that the tertiary γ′-precipitates are dissolving ahead of the crack consistent with the formation of oxides such as alumina and rutile. Our results have implications for alloy design efforts; at any given strength level, both more and less damage-tolerant variants of these alloys can be designed.

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I. INTRODUCTION

BRITTLENESS due to surface-initiated intergranular attack by oxygen is the major limitation to the application of polycrystalline Ni-based superalloys, particularly in the mid-temperature regime of 600 °C to 800 °C.[1] Hence, crack initiation and/or growth can be accelerated under either static or fatigue loading situations, and therefore this phenomenon is of significant importance for high-temperature applications.[2] Alloying has been found to play a role in alleviating this effect, presumably because the chemical reactions which are promoted are then altered in subtle ways.[3] However, these are not fully understood since the mechanism responsible for intergranular crack propagation is yet to be established. Whilst this is the case, it seems unlikely that the chemistry of these alloys has yet been optimized.

Recent work has emphasized the formation of a layered oxide structure consisting of different oxide phases growing ahead of an intergranular crack. This suggests that an oxidation reaction is occurring and that it is not oxygen in its elemental form which is embrittling.[4–6] Thermodynamically less stable oxides consisting of Ni and Co are found at the beginning and center of the crack while more stable oxides consisting of Ti and Al were found closer to the interface of the oxide and metal.[6] In particular, superalloys containing Nb are considered susceptible to environmentally-assisted cracking due to the formation of Nb-rich oxides such as NbO₂ and Nb₂O₅ at the grain boundary ahead of the crack tip.[7,8] But any interpretation is made very much more complex by the interaction of chemistry, microstructure, processing, and the details of the grain boundary reactions which can arise.[9] In addition, damage tolerance arising from microstructure-dependent crack tip shielding and relaxation has to be considered.[10] There are two points which are of overarching importance if progress is to be made towards improved understanding of this effect and hence its alleviation. First, there is a need for systematic experimentation involving the control of the important compositional and microstructural variables such that the effects of others can be isolated, identified, and confirmed. Second, there is a need to better understand the chemical processes occurring at the grain boundaries at the most appropriate lengthscale and with techniques which are capable of providing the most revealing...
information. Scanning transmission electron microscopy/energy-dispersive spectroscopy (STEM-EDS) and high-resolution electron back-scatter diffraction (HR-EBSD) are particularly well suited in this regard, and these are used in the present work.

In this study, three model powder metallurgy (PM) Ni-based superalloys with varying Nb/Ti ratio are studied. Nb partitions to the $\gamma'$-phase and in particular the Al site; it also increases the anti-phase boundary (APB) energy and thus is known to promote strength. Ti behaves similarly. Both are strong oxide formers, but their thermodynamic potentials for doing so differ and therefore one might reasonably expect the Nb/Ti ratio to influence the damage tolerance displayed by these alloys in oxidative environments. Both, slow strain-rate tests and single edge notch tension tests at 700 °C were performed, to elucidate the role of Nb and Ti on the strength, ductility, and toughness displayed in this system. Miniature testpieces of enhanced surface/volume ratio are employed, in order to accelerate the embrittlement effect and thus facilitate its study. A further increase in Nb levels with Nb being substituted for Ti on a 1:1 basis, using the alloys-by-design of approach and deliberately resulting in the same $\gamma'$-precipitate volume fraction. Each ABD-alloy was manufactured by ATI powder metals using a lab-scale version of a commercial powder metallurgy process. Powders for each alloy composition were compacted using hot isostatic pressing (HIP) which involved the application of 100 MPa at 1107 °C. These HIP compacts were isothermally forged at 1100 °C at 0.2 min $^{-1}$ and then given a super-solvus heat treatment conducted at 1170 °C for 2 hours followed by a controlled quench of 1 °C s $^{-1}$ to 1050 °C. Finally, an aging heat treatment at 850 °C was performed for 4 hours. Thermodynamic calculations using Thermocalc were performed to estimate the volume fraction of the $\gamma$-phase and $\gamma'$-phases resulting from the final aging heat treatment at 850 °C (see Table I). Only major phases of interest were included from the thermodynamic database: $\gamma$, $\gamma'$, MC phases, all other phases were suspended. The results are summarized in Table I. The $\gamma'$-solvus temperatures were calculated to be 1157 °C, 1156 °C, and 1154 °C with Thermo-Calc using the Thermotech TTNi8 thermodynamic database for ABD-2, ABD-4, and ABD-6, respectively.

The microstructure of each alloy was assessed using SEM and EBSD characterization. Differences in the grain size were observed (Figure 1(a)) and quantified though a Gaussian curve fitted to the grain size distribution obtained from EBSD analysis, see the final precipitation shearing stress was estimated.
| Alloy | Phase Fraction (Molecular Percent) | Ni (Atomic Percent) | Cr (Atomic Percent) | Co (Atomic Percent) | W (Atomic Percent) | Al (Atomic Percent) | Ti (Atomic Percent) | Ta (Atomic Percent) | Nb (Atomic Percent) | C (Atomic Percent) | B (Atomic Percent) |
|-------|-----------------------------------|---------------------|---------------------|---------------------|--------------------|--------------------|---------------------|---------------------|---------------------|------------------|-------------------|
| ABD-2 nominal | — | 48.5 | 18.9 | 18.5 | 1.0 | 7.9 | 4.3 | 0.7 | 0.0 | 0.13 | 0.05 |
| ABD-2 γ-phase | 51.1 | 36.4 | 33.7 | 25.7 | 1.1 | 2.6 | 0.3 | 0.0 | 0.0 | 0.01 | 0.16 |
| ABD-2 γ′-phase | 48.7 | 63.0 | 3.0 | 10.5 | 0.7 | 13.7 | 8.1 | 1.1 | 0.0 | 0.00 | 0.00 |
| ABD-2 MC-phase | 0.24 | 0.0 | 0.1 | 0.0 | 1.8 | 0.0 | 42.7 | 7.0 | 0.0 | 47.40 | 0.00 |
| ABD-4 nominal | — | 48.5 | 18.6 | 18.6 | 0.9 | 7.9 | 4.0 | 0.6 | 0.4 | 0.01 | 0.10 |
| ABD-4 γ-phase | 52.0 | 36.7 | 33.4 | 25.7 | 1.1 | 2.6 | 0.2 | 0.0 | 0.0 | 0.0 | 0.15 |
| ABD-4 γ′-phase | 47.8 | 63.3 | 2.8 | 10.2 | 0.6 | 13.9 | 7.2 | 1.1 | 0.8 | 0.00 | 0.00 |
| ABD-4 MC-phase | 0.25 | 0.0 | 0.0 | 0.0 | 0.1 | 0.0 | 21.7 | 1.4 | 27.5 | 48.27 | 0.00 |
| ABD-6 nominal | — | 48.1 | 18.9 | 18.8 | 0.9 | 8.1 | 2.9 | 0.6 | 1.2 | 0.12 | 0.11 |
| ABD-6 γ-phase | 52.3 | 36.6 | 33.4 | 25.9 | 1.1 | 2.5 | 0.2 | 0.0 | 0.1 | 0.01 | 0.15 |
| ABD-6 γ′-phase | 47.5 | 63.7 | 2.6 | 9.8 | 0.6 | 14.1 | 5.7 | 1.2 | 2.3 | 0.00 | 0.00 |
| ABD-6 MC-phase | 0.26 | 0.0 | 0.0 | 0.0 | 0.0 | 0.0 | 12.5 | 0.8 | 37.8 | 47.75 | 0.00 |

Phase compositions were calculated with Thermo-Calc using the Thermotech TTNi8 thermodynamic database.
using a model proposed by Galindo-Nava et al.\textsuperscript{[22]} which allows one to calculate the $\gamma'$-precipitation hardening contributions using an estimate mean radius for the secondary and tertiary $\gamma'$-precipitate size, $r_s$ and $r_t$:

$$
\sigma_p = M \frac{\gamma_{APB}}{2b} \left( w_s \frac{l_s}{\Lambda_s} + w_t \frac{l_t}{\Lambda_t} \right),
$$

where $M$ is the Taylor orientation factor; $\gamma_{APB}$ is the anti-phase boundary energy of the $\gamma'$-phase; $b$ is the magnitude of the Burgers vector; $w_s$ and $w_t$ are the particle number fractions of secondary and tertiary $\gamma'$-precipitates; $l_s$ and $l_t$ are the average segment lengths of the leading dislocation acting in the cutting of a secondary and tertiary $\gamma'$-precipitate; $\Lambda_s$ and $\Lambda_t$ are the average effective lengths of the leading dislocations driving the secondary and tertiary $\gamma'$-precipitate cutting. Values for the Taylor orientation factor and the Burgers vector are taken to be 3 and 0.248 nm, respectively. The anti-phase boundary energy of the $\gamma'$-phase at 0K was obtained from theoretical predictions employing DFT and is as follows: $\gamma_{APB}^{\text{ABD-2}} = 0.344$, $\gamma_{APB}^{\text{ABD-4}} = 0.349$, $\gamma_{APB}^{\text{ABD-6}} = 0.36$ Jm$^{-2}$\textsuperscript{[14]}. These values were fitted to the following polynomial function to capture the effect of temperature at 700 °C as proposed by Galindo-Nava et al.\textsuperscript{[22]}: $\gamma_{APB} = \gamma_{APB}^0 - (7 \times 10^{-6} T - 3 \times 10^{-8} T^2)$.

The $\gamma'$-precipitate shearing stress is controlled by strong-pair coupling since both $r_s$ and $r_t$ are greater than the particle radius when the maximum strength is defined by $\tau_m = \mu b^2/2\gamma_{APB}$, with $\mu = E/(2(1 + v))$ and $v = 0.32$. The following analytical expressions were used to capture the effect of temperature $\mu = 87.3 - 9 \times 10^{-4} T - 1.9 \times 10^{-5} T^2$\textsuperscript{[22]}.

The length of the leading dislocation cutting the precipitates is expressed as $l = 2(r_t^2 - (r_m - r_m)^2)^{1/2}$ and the effective length of the leading dislocation driving $\gamma'$-precipitate cutting is given by $\Lambda = \max\{\lambda, L - l\}$, $\lambda$ is the Friedel sampling length which is defined by $\lambda = (2T_{\text{ten}}/r \times \gamma_{APB})$, where $T_{\text{ten}}$ is the dislocation line tension and given by $0.5 \times \mu b^2$ and $L$ is the mean particle spacing $L = (2\pi/3f_\gamma)^{1/2} \times r$.

C. Experimental Assessment of the Mechanical Properties

Tensile testing was conducted at a slow strain-rate of $1 \times 10^{-4}$ s$^{-1}$ at room temperature and at 700 °C using a 5 kN Instron electro-thermal mechanical testing (ETMT) machine. Detailed information relating to the design and operation of the ETMT equipment can be found elsewhere\textsuperscript{[23]}.

Miniaturized specimens of cross section $1 \times 1 \text{ mm}^2$ and a gauge length of 14 mm were
used; the full sample length was 40 mm. Great care was taken with preparation of specimens. These were produced using wire-based electrical discharge machining (EDM), with surfaces subsequently ground to 1200 grit to maintain a consistent surface finish and to remove the recast layer resulting from the EDM process. Consistent with the design of the ETMT machine, specimens were heated using electrical resistance by passing a direct current through the specimen. Temperature measurement during testing was achieved by spot-welding a K-type thermocouple to the central portion of the specimen gauge. A minimum of three tests were performed at each condition, at a constant extension rate consistent with a linear variable displacement transducer (LVDT). The engineering strain was used as a measure of the ductility; engineering strain is defined as the correlation measurements of a thermal paint pattern applied in the central 5 mm of the specimen where the temperature and therefore deformation are uniform\(^{12}\); the engineering stress–strain curves produced were shown to be independent to small changes of \(L_0\).

To quantify the conditional fracture toughness at 700 °C in air, additional ETMT testing was carried out using single edge notch tension specimens. The specimens described above were modified by machining a circular notch on one side to a depth of 150 μm. From this notch, a low cycle fatigue crack was grown at room temperature under load control using an R-ratio of 0.1 and a 2 Hz sinusoidal cycling waveform. A maximum stress level of 95 pct of \(\sigma_y\) was applied. Hereafter the pre-fatigue cracked test pieces were subjected to a constant extension rate of 5 × 10\(^{-4}\) mm s\(^{-1}\) at 700 °C causing the environmentally-assisted growth of the crack. Upon crack growth, tests were interrupted by unloading and simultaneously reducing the temperature to ambient with a cooling rate close to 100 °C s\(^{-1}\). The conditional fracture toughness, \(K_q\), was calculated with conditional load \(P_q\) obtained from the load displacement curve and the measured initial crack length, \(a\), according to the following equation:

\[
K_q = \frac{P_q}{A} \sqrt{a} \times Y,
\]

where \(P_q\) is the maximum load that the specimen could withstand before quasi-brittle crack propagation occurred, and \(A\) is the specimen cross section area. The term \(Y\) is a geometry factor, taken from Reference 24 and is given by

\[
Y = 1.12 - 0.23 \left(\frac{a}{w}\right) + 10.6 \left(\frac{a}{w}\right)^2
- 21.7 \left(\frac{a}{w}\right)^3 + 30.4 \left(\frac{a}{w}\right)^4,
\]

where \(w\) is the specimen width. The determined conditional fracture toughness, \(K_q\), has to be regarded as a geometry-dependent material parameter, due to the amount of crack tip constraint arising due to the miniaturized ETMT tests.

### D. Fractography and Damage Analysis

The microstructure of each alloy both before and after testing was characterized with a Zeiss Merlin scanning electron microscope (SEM). Secondary backscattered electron imaging and HR-EBSD analysis were used. Diffraction patterns were acquired with a Bruker Quantax EBSD system—at a 800 x 600 pixels\(^2\) resolution to provide suitable angular resolution, and saved for subsequent analysis. Using an in-house method to measure the lattice curvature, the distribution of geometrically necessary dislocations (GNDs) was estimated following Wilkinson et al.\(^{25}\).

Fractography was supported by a Zeiss Merlin FEG SEM equipped with an in-column energy selective backscattered (ESB) detector and an Oxford Instruments XMax 150 mm\(^2\) detector for obtaining SEM-EDX elemental maps. Samples for transmission electron microscopy (TEM) were prepared by focused ion beam thinning in a Zeiss Auriga dual beam system. Annular bright-field (ABF) and high angle annular dark field (HAADF) images as well as STEM-EDS elemental maps were acquired in a probe corrected FEI Titan Themis 60-300 microscope equipped with a ChemiSTEM Si-drift EDS-system operated at 300 kV. A probe semi-convergence angle of 17 mrad was used to form the electron probe. The semi-collection angles for ABF and HAADF imaging conditions were chosen to be 8 to 16 and 17 to 73 mrad, respectively.

### III. RESULTS

#### A. Characterization of the Mechanical Properties

Engineering stress–strain curves are shown in Figure 3 at both ambient temperature and 700 °C at a strain-rate of 1 × 10\(^{-4}\) s\(^{-1}\). At room temperature, the alloys showed no significant variation in yield stress, at 935 ± 10 MPa, and ultimate tensile stress (UTS), at 1460 ± 0 MPa; moreover, the tensile ductility showed no dependence on the Nb/Ti ratio and was determined to be 26.7 ± 0.1 pct. At 700 °C all three alloys showed the same 0.2 pct proof stress of 792 ± 3 MPa but the tensile ductility now varied appreciably. The UTS increased with increasing ductility due to the increased amount of work hardening as a function of sample elongation. For tests at 700 °C, ABD-2 showed the least ductility at 4.2 ± 1 pct whilst ABD-4 showed an increase in ductility to 5.4 ± 0.6 pct. The highest tensile ductility was observed for ABD-6 at 7.8 ± 1.1 pct. This is illustrated in Figure 3(b) together with SEM images of cross sections through failed specimens. The tensile properties obtained for each alloy are summarized in Table II. Thus, tensile ductility in the mid-temperature regime increased as a function of the Nb/Ti ratio (see Figure 4(a)) despite the alloys all having the same yield stress. The observed loss in ductility in the mid-temperature regime is consistent with recent observations made on other superalloy systems\(^{26–28}\) and can be explained by the occurrence of intergranular quasi-brittle failure. Crack propagation along the grain boundaries at 700 °C also explains the relatively low conditional fracture.
toughness (see Figure 4(b)). Just like the tensile ductility, toughness increased with increasing Nb/Ti ratio. ABD-2 exhibited the lowest conditional fracture toughness with 
$19.7 \pm 0.5$ MPa$^{\text{ffiffiffiffi}}$ followed by $21.8 \pm 1.2$ and $24.6 \pm 0.4$ MPa$^{\text{ffiffiffiffi}}$ for ABD-4 and 6, respectively. This is summarized in Table III together with the maximum applied load $P_{\text{max}}$ and the estimate pre-crack length, $a$, from which the conditional fracture toughness $K_q$ was calculated using Eq. [4]. The pre-crack and sample geometry were maintained between the tests so that comparisons between alloys are valid.

The yield stress obtained at 700 °C is constant regardless of the apparent changes in Ti/Nb ratio and grain size. To rationalize this, the yield stress was theoretically assessed for each alloy composition including the three dominant strengthening contributions, as described in Section II–B: (1) Hall–Petch strengthening ($\sigma_0$), (2) solid solution strengthening within the $\gamma$-matrix ($\sigma_{ss}$), and (3) resistance to shearing of the secondary and tertiary $\gamma'$-precipitates ($\sigma_p$). The estimated strengthening contributions for each alloy are summarized in Table IV.

The yield strength modeling predictions are in good agreement with the experimental results. Figure 5 shows the yield stress prediction at 700 °C and compares it against the experimental results. By comparing individual strength contributions, one can see that $\gamma'$-precipitate shearing resistance provides the majority of the strength in the mid-temperature regime whilst Hall–Petch strengthening is the lowest component of the overall yield stress. The predicted strength loss with increasing grain size for ABD-6, a stress decrease of 43 MPa, is counteracted by the increase in the $\gamma'$-precipitate shearing stress and $\gamma$-solid solution strengthening resulting in a stress increase of 24 MPa. The APB energy of the $\gamma'$-phase increases with increasing Nb additions thereby increasing its resistance to cutting. The results thus far confirm that for alloy of an equivalent strength, there can be substantial difference in ductility and toughness, resulting from changes in phase chemistry and alloy microstructure, in particular grain size.

B. Characterization of Environmentally-Assisted Failure

The loss in tensile ductility at 700 °C is associated with an increase in the ease of intergranular crack propagation. Intergranular regions on the fracture surface failed perpendicular to the applied load and showed signs of environmental damage. Regions without environmental damage failed at 45 deg to the tensile axis. Figure 6(a) shows a representative secondary electron SEM image of the intergranular fracture surface at 700 °C in ABD-2. The fracture surfaces observed for each alloy showed similar characteristics. Secondary cracks branch off the primary crack which led to failure. Secondary cracks were consistently found along high angle grain boundaries which act as preferred paths for crack propagation (Figures 6(b) and (e)). Moreover, SEM-EDS mapping showed that the cracked grain boundaries are decorated with oxidation products, see Figure 6(c). It is suggested that secondary cracks did not propagate at the time of failure. This could have otherwise lead to subsequent oxidation and have altered the damage ahead of the crack tip.

Primary cracks from interrupted miniaturized single-edge notched crack growth tests in alloy ABD-2 and 6 were chosen for further characterization using high-resolution characterization techniques such as EBSD analysis to provide an estimate of the GND density distribution and STEM-EDS mapping for elemental distribution. All characterization was performed along the crack growth direction normal to the crack face in the center of the testpiece (see Figure 7(a)). Figures 7(b) and (c) show two EBSD inverse pole figure maps with respect to the loading direction on
sections through interrupted primary cracks in ABD-2 and 6, respectively. A transition from transgranular to intergranular cracking can be observed after the first \( \approx 50 \) \( \mu \)m. The transgranular crack growing from the notch is a result of the room temperature fatigue pre-cracking. The fine-grained alloy ABD-2 shows a straight fracture path. In contrast, the crack in the coarse-grained ABD-6 is more tortuous and the opening displacement is reduced. HR-EBSD has been used to calculate an estimate of the GND density distribution with a step size of 450 nm in a 200 \( \times \) 175 \( \mu \)m area within the wake of each crack. In ABD-2, this area showed a low overall GND density. This is observed in Figure 7(d) and highlights the quasi-brittle nature of the intergranular crack propagation. Regions neighboring the intergranular crack show higher GND densities. In addition, there is evidence of a GND storage at the grain boundaries and at triple junctions indicating some amount of deformation at or near the grain boundaries. Figure 7(e) shows that in ABD-6 the GND storage near grain boundaries is less localized suggesting that intragranular deformation is more pronounced. A high GND

| Alloy | Temperature (°C) | YS (MPa) | UTS (MPa) | Tensile Ductility (Percent) |
|-------|-----------------|----------|-----------|----------------------------|
| ABD-2, 4, 6 | 23 | 935 ± 10 | 1460 ± 0 | 26.7 ± 0.1 |
| ABD-2 | 700 | 794 ± 8 | 911 ± 12 | 4.2 ± 1 |
| ABD-4 | 700 | 795 ± 10 | 942 ± 13 | 5.4 ± 0.6 |
| ABD-6 | 700 | 788 ± 14 | 979 ± 5 | 7.8 ± 1.1 |

Mean results and standard deviation (SD) of a minimum of three test repetitions are given.

| Alloy | Yield Stress Prediction (MPa) | \( \sigma_d \) (MPa) | \( \sigma_{ss} \) (MPa) | \( \sigma_p \) (MPa) |
|-------|-------------------------------|-------------------|-------------------|-------------------|
| ABD-2 | 804                          | 117               | 156               | 531               |
| ABD-4 | 795                          | 99                | 160               | 535               |
| ABD-6 | 786                          | 74                | 163               | 548               |

Mean results and SD of a minimum of three test repetitions are given.
density is revealed in a bridging grain which is surrounded by the intergranular crack. Figures 7(f) and (g) show a magnified EBSD and GND map of the intergranular crack tip in ABD-6. The GND density distribution here was calculated with a 100 nm step size showing the accumulation of damage at locations where the grain boundary is serrated and directly ahead of the crack tip. At the tip, the crack is deflected around a smaller grain. In addition, damage accumulation and cracking of a particle, presumably TiN, can be seen within the plastic zone ahead of the crack tip.

To study the nature of the environmental attack leading to the observed intergranular failure, TEM samples were extracted containing a primary crack tip as demonstrated in Figure 8. The Z-contrast HAADF-STEM images shown in Figures 9(a) and (b) illustrate an environmentally-damaged zone ahead of the crack tip. The size of this was found to be independent of alloy composition at a total length of $\approx 770$ nm and a width of $\approx 130$ nm narrowing down into the undamaged grain boundary ahead—while intergranular environmental damage is most pronounced the environmental damage can be observed to penetrate the grains (see Figures 9(c) and (d)). No significant differences in the damage morphology were observed as a function of Nb/Ti ratio. Figures 9(e) and (f) show composite STEM-EDS elemental maps. The overlap of O-K, and Cr-L$_{\alpha}$ X-rays was not found to influence the qualitatively obtained elemental distributions which show the relative elemental distribution integrated over the specimen thickness. More details of the surrounding microstructure such as the tertiary $\gamma'$-precipitates are clearly visible in specimen ABD-2 compared to ABD-6. This effect is related to an increased sample thickness of ABD-6. From these images, a Co and Cr-rich region can be identified mainly in the middle of the damage zone at the $\gamma$-interfaces. Regions which are Ni- and Cr-rich are present close to the damage zone at the $\gamma'$-interfaces. There is evidence suggesting the formation of a distinct Cr-rich oxide layer present in both alloys. In addition, O is observed to penetrate secondary $\gamma'$-precipitates. Internal oxides where found to be Ti and Al-rich. Nb and Ta were found to accumulate in the $\gamma'$-precipitates but was not observed to concentrate in the O-rich regions. Close to the undamaged grain boundary in ABD-6, a Ti and Nb-rich particle, presumably a laves phase, is shown in Figure 9(f). This particle did not exhibit enhanced oxidation damage.

A Ni-rich single phase tertiary $\gamma'$-precipitate free zone is observed in ABD-2 ahead of the environmental damage zone closest to the undamaged grain boundary. This can be observed best in the individual STEM-EDS images in Figures 9(e), 10(b) and (c). Figure 10(b) shows individual STEM-EDS elemental maps and Figure 10(c)
demonstrates the results of a linescan taken across the sub-boundary of the Ni-rich single phase $\gamma'$-phase depleted zone. Striking is the observation that oxygen seems to penetrate a sub-boundary along the Ni-rich region and not the grain boundary. At this sub-boundary, an enrichment in Ti and Al is observed. Ni, Co, and Cr are depleted in this location.

IV. DISCUSSION

The results presented above demonstrate a relationship between the Nb/Ti ratio and the alloys susceptibility to environmentally-assisted cracking at 700 °C. Nonetheless, a number of unresolved issues emerge and these are now considered in this section of the paper.

A. On the Mechanism of Environmentally-Assisted Grain Boundary Embrittlement

The nature of the mechanism of embrittlement in these alloy systems has been the subject of intense interest; one significant issue is the challenge associated with the choice of most appropriate analytical tool for the characterization of damage. Here, STEM-EDS in the TEM has been employed on carefully prepared crack tips; this combination has been particularly insightful. An environmentally-damaged zone penetrating the grain boundary ahead of primary crack tips has been confirmed; within this the formation of a tertiary $\gamma'$-precipitate free-zone is formed which complements the commonly emphasized oxide formation close to the crack tip, which is also observed here. It is this combination of damage which is responsible for intergranular crack propagation which in turn leads to the observed low tensile ductility and low conditional fracture toughness at 700 °C (Figure 4). But which of these is the more important in controlling the loss of ductility?

Particularly pertinent is the observation that the size and order of the reaction products in the environmentally-damaged zone ahead of the crack tip is independent of the Nb/Ti ratio, see Figure 9. The appearance is broadly consistent with the sequence expected from the relative thermodynamic stability of the individual oxidation products, as seen in Figures 9(e) and (f), as reported elsewhere.\[4–6,12,29\] Thus, an Al-rich oxide ($Al_2O_3$) alongside a Ti-rich oxide ($TiO_2$) are found closest to the undamaged grain boundary, resulting from the oxidation of these reactive elements which are found in the $\gamma'$-precipitates, $Ni_3(Al,Ti)$. Therefore, these oxides are associated with extension of the environmentally-damaged zone. This is supported by the observation of a tertiary $\gamma'$-phase depleted zone which is present ahead of the advancing internal oxide as shown in Figures 10(a) and (b). Unlike during the formation of a surface oxide layer, oxygen is readily available at the furthest extent of the oxide intrusion. Because the metal/oxide interfaces are aligned parallel to the direction of oxide growth, they act as short-circuits for oxygen diffusion, see Figure 11. Thus, the damage zone extension during steady-state crack propagation is limited by the dissolution of tertiary $\gamma'$-precipitates and the migration of Al and Ti cations to the oxide/metal interface ahead of the oxide intrusion. For the simple case of $Ni_3Al$ precipitates, the rate controlling reaction is $\gamma' \rightarrow \gamma + Al_2O_3$. The compositional gradient produced
by the depleted zone is assumed to act as an additional driving force for the metal ion diffusion.

It is important to note that the formation of discrete Nb-rich oxides ahead of interrupted primary cracks has not been observed here. Thus, it would appear that any influence of Nb cannot be attributed to its oxidation. This finding is in contrast to the work of Miller & Wei et al. [7,8,30,31] who have studied a number of Nb-containing alloys including IN718. They have suggested that even small additions of Nb increase the susceptibility to environmental-assisted creep crack growth due to the formation of Nb-rich oxides ahead of the crack tip, including NbO₂ and Nb₂O₅. The results presented here indicate that small additions of Nb—up to 2.1 wt pct—are however not detrimental and indeed are advantageous. While the formation of Nb-rich oxides is more favored by thermodynamic equilibrium than TiO₂, Nb is not as abundant as Ti in for example alloy ABD-6 (see Table I) and is believed to stay in solution. This seems likely that the threshold for Nb to have any negative effect is associated with the formation of γ'-precipitates, Ni₃Nb, at the grain boundary. Our findings are in agreement with Telesman et al. [32] who showed that hold time crack growth resistance was insensitive to small changes in the Nb and Ta contents in Alloy 10.

The above suggests that it is the loss of tertiary γ'-precipitates which is detrimental to the localized mechanical properties. This would explain why the tip of the oxide intrusion in ABD-2 does not follow the prior grain boundary (see Figure 10(b)). It nucleates along what is believed to be a newly formed sub-boundary of the single phase tertiary γ'-precipitate depleted zone. It is likely that this interface is formed due to local deformation-induced recrystallization as suggested by Wlodek et al. [33]. The formation of the γ'-phase free zone will be dictated by the tertiary γ'-size and its dissolution kinetics; this is discussed further in Section IV–C.

B. On the Role of Grain Size-Dependent Toughening

Of significance here is the observed increase in grain size with increasing Nb/Ti ratio and its influence on improving the tensile ductility and the conditional fracture toughness. Putting aside for now the issue of why the grain size is influenced by Nb/Ti ratio—this is discussed in the following section—what are the implications of this effect?

While a coarser-grained microstructure resulting from a supersolvus heat treatment has been demonstrated to be beneficial—for example by comparing the behavior of any given alloy to sub-solvus and super-solvus processing [9,32,34–36]—so far, grain size has not been varied independently of the γ'-precipitate size distribution. This gives rise to the question whether grain size independently improves environmentally-assisted crack growth resistance. The yield stress as well as the mean diameter of the secondary and tertiary γ'-precipitates is approximately constant in the present study, so this makes this analysis possible. The HR-EBSD technique has been used for this purpose in order to characterize the deformation in the crack wakes and also the intergranular crack tips. Both the finer-grained ABD-2 and coarser-grained ABD-6 have been studied, in order to deduce any influence of (i) crack bridging as well as (ii) the grain boundary sliding resistance. While the first mechanism will aid resistance to crack propagation along the environmentally-induced damage zone along an embrittled γ'-boundary, an increase in sliding resistance will retard the build-up of a critical strain ahead of the crack tip.

An influence of crack bridging has indeed been observed in the present work, for example in coarser-grained ABD-2, see Figure 7(e), which exhibits significant evidence of this effect in the crack wake. Plastic deformation is also confirmed in GND density maps of this alloy. By way of comparison, grain bridging is not readily apparent in the finer-grained ABD-2, see Figure 7(d). Bridging grains in the wake of the crack is well known to increase the toughness substantially, by reducing the stress intensity at the crack tip. This finding also explains the post-mortem observation present in the literature of unbroken ligaments on the fracture surfaces of testpieces subjected to mid-temperature dwell fatigue. [37–40] The resulting decrease in crack propagation
Fig. 9—HAADF-STEM image of an environmental damage zone ahead of a crack tip in (a) ABD-2 and (b) ABD-6. (c) and (d) show corresponding schematic illustrations highlighting the environmental damage zone ahead of the crack tip. Composite STEM-EDS elemental maps of O with Ni and Co, Al, and Ti as well as Cr are illustrated in (e) and (f). The brightest pixels indicate the highest count areas and dark or black areas indicate little or no X-ray counts. The arrows indicate high X-ray count areas.

Fig. 10—(a) High angle annular dark field STEM image obtained in ABD-2 at the region closest to the undamaged grain boundary shown in Fig. 9(a). Corresponding individual STEM-EDS elemental maps are shown in (b). (c) Linescan obtained across the sub-boundary of the single phase γ′-precipitate depleted zone. A fast Fourier transform filter method was applied to the line profiles.
driving force will contribute towards the observed increase in toughness and ductility as a function of grain size.

It is emphasized that the observed failure mechanism is likely to be closely related to the underlying deformation process. At 700 °C, plasticity is believed to occur through several independent processes including slip-assisted grain boundary sliding mechanism—due to the absence of stress-relieving mechanisms this leads to intergranular fracture and the observed ductility minimum.\[4] The temperature corresponding to the ductility minimum increases with an increased grain size\[41] which explains the observed grain size dependency of the tensile ductility in these alloys, see Figure 3(b). In the finer-grained alloy ABD-2, an increase of GND density at the grain boundaries and grain boundary triple junctions suggests more grain boundary deformation compared to the coarser-grained ABD-6, see Figure 7(d). The increase in grain size as well as the grain boundary serrations inhibit intergranular failure by retarding the formation of local microstresses arising from grain boundary deformation. The observed effects of grain size on ductility and toughness agree with predictions based on a critical crack tip strain fracture model proposed by Floreen and Kane\[42] where stress intensity at the crack tip, \(K\), is proportional to \((\sigma_t \times d)^{1/2}\). A material with a higher resistance to intergranular failure is characterized by a lower strain at the crack tip due to a higher constraint to grain boundary sliding.\[43]

C. On the Relevance to Alloy Design

This work demonstrates that at 700 °C—despite a high yield strength—it is possible to design both more (high Nb/Ti) or less (low Nb/Ti) damage-tolerant alloy variants. For optimum resistance to environmentally-assisted cracking, one should aim for a high yield stress and coarse grain size, in as much as the two can be varied independently of one another. As shown here, resistance to \(\gamma\)-precipitation shearing accounts for the majority of the strength at 700 °C, see Table IV. So what relevance do Ti and Nb play in the present case?

Both Ti and Nb partition strongly to the \(\gamma\)-phase, Ni₃(Al, Ti, Nb), thus influencing the fraction of \(\gamma\)-precipitates and the ease of their shearing via changes in the APB energy.\[14] This is in agreement with the reported experimental results. To achieve a high tensile strength, the \(\gamma\)-phase volume fraction the APB energy has to maximized as suggested by Crudden et al.\[14\] This was done for the alloys designed and studied in this work and can be described by the expression \(M_{\text{strength}} = M \times 1/2 \gamma_{\text{APB}}^{1/2}/b\), where \(M\) is the Taylor factor; \(\gamma\) is the volume fraction of the \(\gamma\)-phase; \(\gamma_{\text{APB}}\) is the anti-phase boundary energy and \(b\) is the Burgers vector. For damage tolerance, it is important to account for the reduction of the Hall–Petch contribution due to a coarse supersolvus microstructure which can be described by \(\sigma_{\text{HP}} = k_y(D_{\text{sub}}^{-1/2} - D_{\text{super}}^{-1/2})\), where \(k_y\) is the Hall–Petch coefficient; \(D_{\text{sub}}\) and \(D_{\text{super}}\) denote the grain size resulting from a sub-solvus or super-solvus heat treatment. This impact of this is illustrated in Figure 12(a) where strength and ductility for a variety of polycrystalline Ni-based superalloys are plotted. The trade-off between strength and tensile ductility can therefore be described by \(M_{\text{strength}} = M \times 1/2 \gamma_{\text{APB}}^{1/2}/b - \sigma_{\text{HP}}\), where a grain size-dependent strength reduction is counteracted by a degree of composition-dependent strengthening.

In addition to grain size, the observed dissolution of the tertiary \(\gamma\)-precipitates during the extension of the environmental damage zone during continuous crack propagation suggests that both size and composition of the tertiary \(\gamma\)-precipitates influence the susceptibility to embrittlement. Thus, an increase in the mean diameter would result in slower dissolution. This is in agreement with Telesman et al.\[45\] and Li et al.\[9\] who have shown that an increase in tertiary \(\gamma\)-precipitate size is beneficial during dwell fatigue loading. The composition of the tertiary \(\gamma\)-precipitates is important since the rate of dissolution and the availability of oxidizing elements at the furthest end of the damage zone will be composition dependent. A more stable tertiary \(\gamma\)-precipitate population ahead of the environmental damage zone will exhibit slower \(\gamma\)-phase dissolution kinetics and therefore a slower rate of oxide formation. Collins et al.\[17\] demonstrated that the Nb present in ABD-6 slows down the formation and dissolution kinetics of the \(\gamma\)-phase compared to ABD-2. This is in agreement which the results presented here.

A last point relates to the reason why the Nb/Ti ratio alters the grain size in this system. It seems likely that this is due to the MC carbide composition, the temperature-dependence of the stability of this phase during forging/subsequent heat treatment and its influence on recrystallization and grain growth. A number of studies have observed the effect of MC carbide composition on the grain size during the processing of polycrystalline Ni-based superalloys and specifically the pinning of grain boundaries by them.\[46,47\] A change in the MC carbide composition from Ti to Nb-rich MC carbides is apparent in the thermodynamic predictions of Table I.
The difference in carbide stability at 1170 °C is also seen in the carbide Ellingham diagram of Figure 12(b). Titanium is known to stabilize MC and hence retard its coarsening and dissolution. This is likely to suppress grain growth at lower Nb/Ti ratio in for example alloy ABD-2. But accounting for the influence of alloy composition on grain size development during processing is a significant issue which should be the topic of future work.

V. SUMMARY AND CONCLUSIONS

In this work, three high-strength Ni-based superalloys of varying Nb/Ti ratio have been studied. The following specific conclusions can be drawn:

(1) Differences in the Nb/Ti ratio did not alter the yield stress of these alloys either at ambient conditions—at which it is ~ 935 MPa—or at 700 °C where it is reduced to ~ 792 MPa. This occurs despite an influence of the increasing Nb/Ti ratio which coarsens the grain size from 18 to 45 μm; the mean diameter of both secondary and tertiary γ’-precipitates is independent of Nb/Ti ratio at 230 and 40 nm, respectively.

(2) The above implies that any loss in Hall–Petch strengthening due to the increase in grain size as the Nb/Ti ratio increases is offset by an increase in γ'-matrix solid solution hardening and/or an increased resistance to γ'-precipitate shearing.

Analysis via theoretical modeling has been used to rationalist the increments in strengthening due to these effects.

(3) In contrast, the tensile ductility at 700 °C is influenced substantially by the Nb/Ti ratio due to an effect on the susceptibility to quasi-brittle intergranular cracking. Tensile ductility is no longer alloy independent and is approximately doubled from 4.2 pct for the smallest Nb/Ti ratio examined (zero) to 7.8 pct for the largest (0.43). Consistent with this, the elevated temperature conditional fracture toughness calculated from interrupted single edge notched tests increases from 19.7 to 24.6 MPa√m.

(4) Characterization of interrupted primary crack tips revealed sub-micron sized oxide intrusions which penetrate the grain boundaries. The size of the intrusions is found to be independent of the Nb/Ti ratio. However, the oxide intrusions contain different phases present in an order consistent with thermodynamics. The internal oxidation of reactive elements found in γ'-precipitates, Al2O3 and TiO2, are responsible for extending the environmental damage zone during steady-state crack propagation. Not observed are Nb-rich oxides such as NbO2.

(5) Whilst all alloys investigated failed at 700 °C in a quasi-brittle fashion due to environmentally-assisted cracking, a coarse grain size increases both the tensile ductility and the fracture toughness. Crack bridging, crack deflection, and an increased
resistance to grain boundary sliding play roles in this. In addition, evidence is presented which confirms that the tertiary γ’-precipitate size and composition affect the susceptibility to environmentally-assisted cracking, since the dissolution of the tertiary γ’-precipitates ahead of the crack is required to form Al2O3 and TiO2 at undamaged grain boundaries.

(6) The present study has implications for the design of damage-tolerant high-strength Ni-based superalloys. At any given strength level, our results demonstrate clearly that more or less damage-tolerant variants can be designed by subtle changes in alloy composition; for optimum resistance to environmentally-assisted cracking one should aim for a high yield stress and coarse grain size, in as much as the two can be varied independent of one another. The balance of the elements Nb and Ti needs to be chosen carefully in this regard.

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REFERENCES

1. R. Jiang, D.J. Bull, D. Proprentner, B. Shollock, and P.A.S. Reed: Int. J. Fatigue, 2017, vol. 99, pp. 175–86, https://doi.org/10.1016/j.ijfatigue.2017.03.003.
2. T.M. Pollock and S. Tin: J. Propuls. Power, 2006, vol. 22, pp. 361–74, https://doi.org/10.2514/1.18239.
3. S. Cruchley, H. Evans, and M. Taylor: Mater. High Temp., 2016, vol. 3409, pp. 1–11, https://doi.org/10.1007/978-0-367-1171952.
4. A.A.N. Németh, D.J. Crudden, D.E.J. Armstrong, D.M. Collins, K. Li, A.J. Wilkinson, C.R.M. Grovenor, and R.C. Reed: Acta Mater., 2017, vol. 126, pp. 361–71, https://doi.org/10.1016/j.actamat.2016.12.039.
5. H.S. Kitaguchi, M.P. Moody, H.Y. Li, H.E. Evans, M.C. Hardy, and S. Lozano-Perez: Scripta Mater., 2015, vol. 97, pp. 41–44, https://doi.org/10.1016/j.scriptamat.2014.10.025.
6. L. Viskari, M. Hörnqvist, K.L. Moore, Y. Cao, and K. Stiller: Acta Mater., 2013, vol. 61, pp. 3630–39, https://doi.org/10.1016/j.actamat.2013.02.050.
7. R.P. Wei, C.F. Miller, Z. Huang, G.W. Simmons, and D.G. Helmick: Eng. Fract. Mech., 2009, vol. 76, pp. 715–27, https://doi.org/10.1016/j.engfracmech.2008.09.003.
8. G. Németh, D.J. Crudden, D.M. Collins, D.E.J. Armstrong, and R.C. Reed: Superalloys, Wiley, Hoboken, 2016, pp. 549–60.
9. M. Gao, D.J. Dwyer, and R.P. Wei: Scripta Metall. Mater., 1995, vol. 32, pp. 1169–74, https://doi.org/10.1016/0956-716X(95)00120-K.
10. H.Y. Li, J.F. Sun, M.C. Hardy, H.E. Evans, S.J. Williams, T.J.A. Doel, and P. 2015: Acta Mater., 2015, vol. 90, pp. 355–69, https://doi.org/10.1016/j.actamat.2015.02.023.
11. J. Telesman, T.P. Gabb, and L.J. Ghosn: Superalloys, Wiley, Hoboken, 2016, pp. 801–10.
12. R.C. Reed, T. Tao, and N. Warnken: Acta Mater., 2009, vol. 57, pp. 5898–13, https://doi.org/10.1016/j.actamat.2009.08.018.
13. D.J. Crudden, A. Mottura, N. Warnken, B. Raesinisa, and R.C. Reed: Acta Mater., 2014, vol. 75, pp. 356–70, https://doi.org/10.1016/j.actamat.2014.04.075.
14. J.O. Andersson, T. Helsander, L. Höglund, P. Shi, and B. Sundman: Calphad, 2002, vol. 26, pp. 273–312, https://doi.org/10.1016/S0364-5916(02)00037-8.
15. C.A. Schneider, W.S. Rasband, and K.W. Eliceiri: Nat. Methods, 2012, vol. 9, pp. 671–75, https://doi.org/10.1038/nmeth.2009.
16. D.M. Collins, D.J. Crudden, E. Alaberti, T. Connolley, and R.C. Reed: Acta Mater., 2015, vol. 94, pp. 244–56, https://doi.org/10.1016/j.actamat.2015.04.046.
17. F. Wallow and E. Nembach: Scripta Mater., 1996, vol. 34, pp. 499–505, https://doi.org/10.1016/S0956-716X(95)00548-4.
18. I.A. Gypen and A. Deruyttere: J. Mater. Sci., 1977, vol. 12 (5), pp. 1028–33, https://doi.org/10.1007/BF00540987.
19. H.A. Roth, C.L. Davis, and R.C. Thomson: Metall. Mater. Trans. A, 1997, vol. 28A, pp. 1329–35, https://doi.org/10.1007/s11661-997-0268-2.
20. Y. Mishima, S. Ochiai, N. Hamao, M. Yoshogawa, and T. Suzuki: Trans. Jpn. Inst. Metals, 1986, vol. 27, pp. 656–64, https://doi.org/10.2320/matertrans1960.27.656.
21. E.I. Galindo-Nava, L.D. Connor, and C.M.F. Rae: Acta Mater., 2015, vol. 98, pp. 377–90, https://doi.org/10.1016/j.actamat.2015.07.048.
22. B. Roebuck, D.C. Cox, and R.C. Reed: Superalloys, Wiley, Hoboken, 2004, pp. 523–28.
23. D.P. Rooke and D.J. Cartwright: Eng. Fract. Mech., 1976, vol. 8, pp. 567–73, https://doi.org/10.1016/0013-7944(76)90010-2.
24. P.D. Littlewood, T.B. Britton, and A.J. Wilkinson: Acta Mater., 2011, vol. 59, pp. 6489–6500, https://doi.org/10.1016/j.actamat.2011.07.016.
25. M.C. Rezende, L.S. Araújo, S.B. Gabriel, J. Dille, and L.H. de Almeida: J. Alloys Compd., 2015, vol. 643, pp. 256–59, https://doi.org/10.1016/j.jallcom.2014.12.279.
26. Gabb, T.P., Telesman, J., Banki, A., and McDevitt, E.: Proceedings of the Symposium on Superalloy 718 and Derivatives, 2014, pp. 697–712.
27. M. Calmunger, G. Chai, S. Johansson, and J. Moverare: Theor. Appl. Mech. Lett., 2014, vol. 4, pp. 010041–46, https://doi.org/10.1063/2.1400414.
28. H.S. Kitaguchi, H.Y. Li, H.E. Evans, R.G. Ding, I.P. Jones, G. Baxter, and P. Bowen: Acta Mater., 2013, vol. 61, pp. 1968–81, https://doi.org/10.1016/j.actamat.2012.12.017.
29. C.F. Miller, G.W. Simmons, and R.P. Wei: Scripta Mater., 2000, vol. 42, pp. 227–32, https://doi.org/10.1016/S1359-6462(99)00336-X.
30. C.F. Miller, G.W. Simmons, and R.P. Wei: Scripta Mater., 2003, vol. 48, pp. 103–08, https://doi.org/10.1016/S1359-6462(02)00355-X.
31. J. Telesman, P. Kanta, J. Gayda, P.J. Bonacuse, and A. Prescenzzi: Superalloys, Wiley, Hoboken, 2004, pp. 215–24.
32. S.T. Wloch: Trans. Metall. Soc. AIME, 1964, vol. 230, pp. 1078–90.

METALLURGICAL AND MATERIALS TRANSACTIONS A
34. H.T. Pang and P.A.S. Reed: Mater. Sci. Eng. A, 2007, vol. 448, pp. 67–79, https://doi.org/10.1016/j.msea.2006.11.016.
35. S. Everitt, M.J. Starink, and P.A.S. Reed: Superalloys, Wiley, Hoboken, 2008, pp. 714–50.
36. R. Jiang, S. Everitt, M. Lewandowski, N. Gao, and P.A.S. Reed: Int. J. Fatigue, 2014, vol. 62, pp. 217–27, https://doi.org/10.1016/j.ijfatigue.2013.07.014.
37. D. Gustafsson and E. Lundström: Int. J. Fatigue, 2013, vol. 48, pp. 178–86, https://doi.org/10.1016/j.ijfatigue.2012.10.018.
38. M. Hörmqvist, L. Viskari, K.L. Moore, and K. Stiller: Mater. Sci. Eng. A, 2014, vol. 609, pp. 131–40, https://doi.org/10.1016/j.msea.2014.04.102.
39. U. Krupp, K. Wackermann, and H.-J. Christ: Magnus Hörmqvist Colliander, and Krystyna Stiller: OxidMet., 2017, vol. 88, pp. 3–14, https://doi.org/10.1007/s11085-016-9707-z.
40. R. Jiang and P.A.S. Reed: Mater. Sci. Technol., 2016, vol. 32, pp. 401–06, https://doi.org/10.1080/02670836.2016.1148227.
41. S.L. Mannan, K.G. Samuel, and P. Rodriguez: Mater. Sci. Eng., 1985, vol. 68, pp. 143–49, https://doi.org/10.1016/0025-5416(85)90403-3.
42. S. Floreen and R.H. Kane: Metall. Mater. Trans. A, 1976, vol. 7A, pp. 1157–60, https://doi.org/10.1007/BF02656598.
43. K. Maciejewski and H. Ghonem: Mater. Sci. Eng. A, 2013, vol. 560, pp. 439–49, https://doi.org/10.1016/j.msea.2012.09.088.
44. S.R. Shatynski: Oxid. Metals, 1979, vol. 13, pp. 105–18, https://doi.org/10.1007/BF00611975.
45. J. Telesman, T.P. Gabb, A. Garg, P. Bonacuse, and J. Gayda: Superalloys, Wiley, Hoboken, 2008, pp. 807–16.
46. G.H. Gessinger and M.J. Bomford: Int. Metall. Rev., 1974, vol. 19, pp. 51–76, https://doi.org/10.1179/imtr.1974.19.1.51.
47. B. Piekarski: Mater. Charact., 2001, vol. 47, pp. 181–86, https://doi.org/10.1016/S1044-5803(01)00166-8.
48. M.F. Ashby and K.E. Easterling: Acta Metall., 1982, vol. 30, pp. 1969–78, https://doi.org/10.1016/0001-6160(82)90100-6.