ABSTRACT

The Ni-33Al-33Cr-1Mo eutectic has been directionally solidify by a modified Bridgeman technique at growth rates ranging from 7.6 to 508 mm/h to produce grain/cellular microstructures containing alternating plates of NiAl and Cr alloyed with Mo. The grains had sharp boundaries for slower growth rates (≤ 12.7 mm/h), while faster growth rates (≥ 25.4 mm/h) lead to cells bounded by intercellular regions. Compressive testing at 1300 K indicated that alloys DS’ed at rates between 25.4 to 254 mm/h possessed the best strengths which exceed that for the as-cast alloy.

INTRODUCTION

Directional solidification (DS) of NiAl-X systems has shown promise for the simultaneous improvement of elevated temperature strength and room temperature toughness [1-4]. In general it was believed that these benefits could only be possible when the structure was perfectly aligned and fault free. Unfortunately such ideal microstructures tend to demand very slow growth rates which would be impracticable for commercialization. To determine if faster growth rates could produce materials with acceptable elevated temperature strength properties, the eutectic system Ni-33Al-33Cr-1Mo (at %) has been directionally solidified at rates ranging from 7.6 to 508 mm/h. This system was chosen since it forms a lamellar eutectic microstructure [1-3] comprised of NiAl and Cr alloyed with Mo {Cr(Mo)} which has demonstrated a room temperature toughness of 17.3 MPa·m [3]. This paper presents the alloy chemistry, microstructure and 1300 K compressive behavior for both as-cast and directionally solidified materials. Compression was utilized to determine mechanical properties because of the ease of machining and testing specimens and the measurement of identical 1300 K compressive and tensile flow stress - strain rate properties in several DS’ed NiAl-X systems [5,6].

EXPERIMENTAL PROCEDURES

A detailed discussion of the techniques employed to produce directionally solidified rods of 33Ni-33Al-33Cr-1Mo is presented in refs. 4 and 6. In short, 19 mm diameter as-cast alloy bars were prepared by induction melting in Al2O3 crucibles and casting into a copper chill mold. Such bars were then placed in high purity alumina open-ended tubes for directional solidification in a modified Bridgeman apparatus under flowing high purity argon. Preferential solidification was accomplished by pulling the tube through a hole in a fixed position water-cooled copper baffle which yielded thermal gradients at the liquid/solid interface of about 8-10 K/mm. A total of seven DS’ed rods were produced at growth rates of 7.6, 12.7, 25.4, 50.8, 127, 254 and 508 mm/h. Samples for chemistry and metallurgy examination were taken from each as-cast bar as well as the aligned region of each DS’ed rod. Chemical analysis to determine both major and minor solute metallic elements was performed by an inductively coupled plasma (ICP)
technique. The concentrations of nitrogen and oxygen were determined by an inert gas fusion method, while the carbon level was measured by the combustion extraction method. Transverse and longitudinal sections of selected as-cast bars and all seven DS’ed rods were metallographically prepared and examined by light optical techniques.

Parallelepiped compression samples 8 x 4 x 4 mm in size with the long axis parallel to the casting or DS growth direction were electrodischarge machined from several as-cast bars and the aligned region of each DS’ed rod. Both constant load and constant velocity compressive testing was undertaken in air at 1300 K. Lever arm test machines were utilized for constant load compressive creep experiments, where deformation was determined as a function of time by measuring the relative positions of the ceramic push bars applying the load to the specimen. Constant velocity tests were conducted in a universal machine at crosshead rates ranging from $1.7 \times 10^{-2}$ to $1.7 \times 10^{-6}$ mm/s. All the acquired test data were normalized to the final specimen length, and true stresses, strains and strain rates were determined with the assumption of constant volume.

RESULTS AND DISCUSSION

Alloy Composition -- The average, maximum and minimum values and the standard deviation for Al, Cr and Mo are reported for the as-cast bars and directionally solidified rods in Table 1.

| Table 1 Composition of as-cast and directionally solidified Ni-33Al-31Cr-1 Mo alloys |
|---------------------------------|-------------------------------|-------------------------------|-------------------------------|
|                                 | Aluminum, at. % | Chromium, at. % | Molybdenum, at. % |
| As-Cast                      | 33.00 | 33.26 | 32.43 | 0.28 | 33.39 | 33.56 | 33.29 | 0.10 | 1.03 | 1.05 | 1.02 | 0.01 |
| DS’ed                         | 32.85 | 33.71 | 31.21 | 0.68 | 33.47 | 34.90 | 32.88 | 0.54 | 1.02 | 1.04 | 0.99 | 0.016 |

The alloys also contained (at. %) about 0.009Cu, 0.010Fe, 0.005Si, 0.07C, 0.002N, 0.03O and 0.001S as impurities. The results in Table 1 indicate the control of the alloy chemistry for all seven as-cast bars and DS’ed rods was quite good, and the average values are close to the intended Ni-33Al-31Cr-1Mo composition. While there is little change between the Mo levels in the as-cast and DS’ed alloys, surprisingly directionally solidification appeared to increase the difference between the maximum and minimum values for both Al and Cr, and thus Ni, over those measured in the as-cast bars. Such changes are, in turn, reflected by the larger standard deviations for the DS’ed rods.

Alloy Microstructure -- Photomicrographs illustrating the transverse structure of as-cast and DS’ed Ni-33Al-33Cr-1Mo are presented in Fig. 1. Relatively rapid solidification during casting into a Cu-chill mold produced NiAl dendrites scattered among grains containing both NiAl and Cr(Mo) (Fig. 1(a)). This can be compared to the regions (grains) of parallel Cr(Mo) and NiAl plates delineated by sharp boundaries (Fig. 1(b)) after DS’ing at 12.7 mm/h; cells enclosed by relatively thick intercellular regions where each cell contains lamella in a radial pattern (Fig. 1(c)) after DS’ing at 127 mm/h; and cells containing (Cr,Mo) fibers which are surrounded by a thick, coarse structured intercellular border after DS’ing at 508 mm/h. The general appearance of the transverse microstructure in Fig. 1(c) is typical of that found after DS’ing between 50.8 to
254 mm/h. The structure after the slowest growth rate (7.6 mm/h) is similar to that shown in Fig.
1(b) except that occasional cells are partially composed of Cr(Mo) fibers.

In spite of some difference in chemistry among all the as-cast bars and DS’ed rods (Table 
1), no third phases were found nor was there any evidence of NiAl or Cr(Mo) dendrites in the 
DS’ed regions. Overall the microstructures of the present Ni-33Al-33Cr-1Mo rods as a function 
of DS rate are similar to those found in Ni-33Al-31Cr-3Mo [4,7] as a function of growth rate. 
The structure in Fig. 1(b) is also in agreement with that for NiAl-32.4Cr-1Mo grown at 12.7 
mm/h by Cline, et al. [8] utilizing a Bridgeman technique. On the other hand Yang et al. [3] 
were able to maintain the sharp boundary cells comprised of the parallel plate type of 
microstructure (Fig. 1(b)) at growth rates of both 50 and 100 mm/h in Ni-33Al-33Cr-1Mo 
through an Edge-defined Film-fed Growth method.

1300 K Compressive Properties -- Examples of the compressive stress - strain curves obtained 
from constant velocity testing and the creep curves measured under constant load conditions are 
given in Fig. 2. The stress - strain curves (Fig. 2(a-c)) indicated that all materials underwent 
work hardening over the first one percent stain followed by continued flow at a more or less 
constant stress. As indicated by the two pairs of filled and open symbols in Fig. 2(b), testing 
under nearly identical conditions produced essentially the same stress-strain curves. Overall the 
constant velocity data (Figs. 2(a-c)) illustrate that 1300 K strength of the alloys decreases as the 
imposed deformation rate decreases, and at slower strain rates the as-cast material (Fig. 2(a)) is 
weaker than the DS’ed alloys (Fig. 2(b,c)). The creep curves for all the alloys (Fig. 2(d)) 
displayed normal behavior with work hardening during primary creep followed by steady state 
flow. This figure also illustrates the advantage of directional solidification, as like amounts of 
creep strain were accumulated over ~600 ks in spite of the much higher stresses on the two 
DS’ed alloys than on the as-cast material.

The 1300 K plastic flow stress - strain rate properties of as- cast and DS’ed Ni-33Al-
33Cr-1Mo are presented in Fig.3, where flow strength was taken as the stress at 1 % from the 
constant velocity test results (Fig. 2(a-c)) and the average stress over the steady state regime 
from the creep curves (Fig. 2(d)). The results from two as-cast Ni-33Al-33Cr-1Mo bars are 
presented in Fig. 3(a), and they illustrate that both bars possessed alike strengths. While all the 
flow stress - strain rate data (Fig. 3(b)) for the seven directionally solidified rods falls within a 
well defined band, visual examination of the results suggests that the properties of slowest (7.6 
mm/h) and fastest (508 mm/h) DS’ed rods were inferior to Ni-33Al-33Cr-1Mo DS’ed at 
intermediate growth rates. This contention was statistically tested utilizing an exponential stress 
law in combination with a dummy variable, where it was verified that 1300 K deformation 
characteristics of the rods DS’ed between 25.4 and 254 mm/h were equivalent. This group of 4 
rods was statistically superior in strength compared to that of the 7.6 and 508 mm/h materials 
and marginally stronger than the rod DS’ed at 12.7 mm/h.

The 1300 K behavior of the best DS’ed rods of Ni-33Al-33Cr-1Mo are compared to the 
as-cast alloy in Fig. 4(a). This figure also illustrates the linear regression fits of the flow stress 
(σ in MPa) - strain rate (ε in s⁻¹) data for both sets of material. Because of the log-log format of 
Fig. 4(a), the exponential fit for the DS’ed alloys, \( \dot{\varepsilon} = 3.13 \times 10^{-10} \exp(0.051\sigma) \), shows 
curvature; whereas the power law description of the as cast alloy, \( \dot{\varepsilon} = 1.61 \times 10^{-18} \sigma^{6.00} \), is a 
straight line. Taken together the data show that directional solidification yields a very positive 
(~2x) strength advantage over simple casting at strain rates < 10⁻⁵ s⁻¹, but there is no advantage at 
faster deformation rates.
As the 1300 K properties of DS’ed Ni-33Al-33Cr-1Mo are not dependent on growth rates ranging from 25.4 to 254 mm/h, changes in microstructural parameters, such as a refinement in interlamellar spacing, cell diameters and intercellular regions, are either unimportant or act in a manner to counterbalance each other. In comparison to the previous study by Yang, et al. [3], structure could be important since testing of their planar eutectic Ni-33Al-33Cr-1Mo at $\dot{\varepsilon} \sim 5 \times 10^{-4} \text{s}^{-1}$ resulted in ultimate tensile strengths (UTS) of 420 MPa at 1255 K and 348 MPa at 1366 K. Linear extrapolation of these results to 1300 K suggests a UTS of about 390 MPa which is approximate 100 MPa greater than the best flow stress measured in the current study (Fig. 3(b)). While a potential for improvement might exist, the curves in Fig. 4(b) reveal that the 1300 K slow strain rate compressive properties of the current DS’ed Ni-33Al-33Cr-1Mo are slightly better than those of DS’ed NiAl-34Cr and equivalent to those of DS’ed Ni-33Al-28Cr-6Mo [2].

In summary, we were able to reproducibly melt and cast Ni-33Al-33Cr-1Mo and directionally solidify this NiAl + Cr(Mo) two phase eutectic by the Bridgeman technique at growth rates ranging from 7.6 to 508 mm/h. Compressive testing at 1300 K indicated that alloys DS’ed at rates between 25.4 to 254 mm/h possessed the best strengths; hence it might be possible to produce materials with acceptable elevated temperature strength at a reasonably fast DS rate. Future work will concentrate on the 1200 and 1400 K compressive behavior to determine if the same dependency on growth rate is maintained; additionally the room temperature toughness of as-cast and DS’ed alloys will be measured to determine if an optimum combination of properties can be achieved.

SUMMARY OF RESULTS

Seven bars of Ni-33Al-33Cr-1Mo have been directionally solidified at rates ranging from 7.6 to 508 mm/h which generally produced alternating plates of NiAl and (Cr,Mo) in grains with sharp boundaries ($\leq 12.7 \text{ mm/h}$) or cells surrounded by intercellular regions ($\geq 25.4 \text{ mm/h}$). Compressive testing at 1300 K indicated that rods DS’ed at rates ranging from 25.4 to 254 mm/h had the best strength which, in turn, was substantially better than the properties of the as-cast alloy.

REFERENCES

Figure 1. Transverse microstructure of as-cast and directionally solidified NiAl-(Cr,Mo) eutectics. (a) As-cast Ni-33.1Al-33.3Cr-1.03Mo, (b) Ni-32.16Al-34.06Cr-1.03Mo grown at 12.7 mm/h, (c) Ni-32.59Al-33.40Cr-1.04Mo grown at 127 mm/h and Ni-32.86-33.30-1.03Mo grown at 508 mm/h. NiAl is the dark phase and (Cr,Mo) is the bright phase.

Figure 2. True compressive stress-strain curves as a function of strain rate from 1300 K constant velocity testing of Ni-33Al-33Cr-1Mo: (a) As-cast, (b) DS’ed at 25.4 mm/h and (c) DS’ed at 508 mm/h; and true compressive creep curves from constant engineering stress creep testing of three forms of Ni-33Al-33Cr-1Mo: As-cast at 40 MPa, DS’ed at 25.4 mm/h at 70 MPa and DS’ed at 254 mm/h at 75 and 102 MPa.

Figure 3. True 1300 K compressive flow stress-strain rate behavior for Ni-33Al-33Cr-1Mo. (a) two as-cast bars and (b) seven DS’ed rods as a function of growth rate.

Figure 4. Comparison of the 1300 K strength properties of the strongest DS’ed Ni-33Al-33Cr-1Mo rods to those for the (a) As-cast alloy and (b) DS’ed NiAl-Cr(Mo) alloys [2].
Figure 2

(True compressive strain, %)

(a) True compressive stress, MPa

(b) True compressive stress, MPa

(c) True compressive stress, MPa

(d) True compressive strain, %

Legend:
- • 2.7x10^{-3}
- □ 2.5x10^{-4}
- ▲ 2.2x10^{-5}
- ▲ 1.7x10^{-6}
- ○ 3.1x10^{-7}

Non DS'ed
- 254 mm/h
- at 102 MPa
- at 75 MPa
- at 40 MPa

DS'ed 25.4 mm/h
- at 71 MPa
- 62 MPa

Time, ks

Whittenberger 1 Opt!1 00% from author's electronic file
True compressive stress, MPa

True compressive strain rate, s⁻¹

Bar

13/98

11/98

True compressive stress, MPa

True compressive strain rate, s⁻¹

7.6

12.7

25.4

50.8

127

254

508

Figure 3
Figure 4

(a) True compressive stress, MPa vs. True compressive strain rate, s\(^{-1}\)

- ▲ 25.4
- ○ 50.8
- ▲ 127
- ▼ 254
- ◆ As-cast

(b) True compressive strain rate, s\(^{-1}\) vs. True compressive stress, MPa

- Ni-33Al-33Cr-1Mo
- NiAl-34Cr
- NiAl-33Al-28Cr-6Mo

E-12470 Whittenberger 10pt/100% from author's electronic file