HIGH TEMPERATURE STRENGTH OF REFRACTORY COMPLEX CONCENTRATED ALLOYS

O.N. Senkov^{1,*}, S. Gorsse² and D.B. Miracle¹

¹ Air Force Research Laboratory, Materials and Manufacturing Directorate, Wright-Patterson AFB, OH 45433, USA

² Université de Bordeaux, CNRS, Bordeaux INP, ICMCB, UMR 5026, F-33600 Pessac, France

* Corresponding author. Email: oleg.senkov.ctr@us.af.mil

ABSTRACT

Thermodynamic and mechanical properties of 15 single-phase and 11 multi-phase refractory complex concentrated alloys (RCCAs) are reported. Using the CALPHAD approach, phase diagrams for these alloys are calculated to identify the solidus (melting, T_m) temperatures and volume fractions of secondary phases. Correlations were identified between the strength drops at 1000°C and 1200°C and the alloy compositions, room temperature properties, melting temperatures and volume fractions of secondary phases. The influence of alloy density on the temperature dependence of specific yield strength was also explored. The conducted analysis suggests that the loss of high-temperature strength of single-phase BCC RCCAs is related to the activation of diffusion-controlled deformation mechanisms, which occurs at $T \geq 0.6 T_m$, so that the alloys with higher T_m retain their strength to higher temperatures. On the other hand, a rapid decrease in strength of multi-phase RCCAs with increasing temperature above 1000°C is probably due to dissolution of secondary phases.

Keywords: Refractory alloys; mechanical properties; high temperature strength; CALPHAD.

1 INTRODUCTION

Refractory complex concentrated alloys (RCCAs) have recently brought much attention as promising candidates for high-temperature structural applications, beyond Ni-based superalloys [1-3]. An RCCA, which is also called refractory multi-principal element alloy (RMPEA), consists of three or more principal elements and it may also contain non-principal elements, which are generally added to further improve/optimize properties of the principal composition. Refractory high entropy alloys (RHEAs), which, by definition, contain five or more principal elements with concentrations of each principal element between 5 and 35 at%, are a sub-class of RCCAs. While RHEAs, because of their high configurational entropy, are sometimes considered to be single-phase, disordered solid solution alloys, RCCAs expand this vast range of new alloys by including multi-phase alloys. RHEAs were first introduced in 2010 [4] and immediately attracted attention due to their ability to retain high strength at temperatures beyond 1000°C, and in some cases up to 1600°C [1, 5]. In the following years the number of newly reported RCCAs has increased steadily, exceeding 180 by the end of 2018. An extensive review of these alloys and their properties has recently been given [1, 2, 6, 7].

It is well known that metallic alloys generally show a considerable decrease in strength at temperatures above about $0.6T_m$, where T_m is the absolute melting point. Increasing T_m may therefore increase the maximum operating temperature. Thus refractory metals and alloys with extremely high melting points can be potential candidates for applications at temperatures significantly above Ni-based superalloys. High brittle-to-ductile transition temperature, inferior oxidation resistance and high density limit the applications of many conventional refractory alloys based on a single element. The RCCA concept provides much higher flexibility for the selection of alloy compositions for improved properties. In particular, RCCAs with reduced densities (down to 5.5 g/cm³), considerably improved oxidation resistance and improved room temperature ductility and processability have been identified [1].

More than half of the reported RCCAs are single-phase BCC structures. The matrix phase of multi-phase RCCAs has a BCC or B2 crystal structure and the secondary phases are primarily BCC, B2, Laves, silicides or their combination [1]. Due to the large variations in the compositions, phase content, microstructures and preparation/processing methods, reported RCCAs show quite a large range of mechanical properties [1, 2, 6]. For example, the yield strengths of reported RCCAs fall in the range of 400 MPa to 2612 MPa at 25°C, 40 MPa to 935

MPa at 1000°C and 24 MPa to 979 MPa at 1200°C [6]. Multi-phase RCCAs are generally stronger at room temperature (RT) but they lose their strength more rapidly with increasing temperature as compared to single-phase RCCAs [1]. As a result, single-phase RCCAs become stronger than many multi-phase RCCAs at $T \geq 1000^\circ\text{C}$. For each of these two alloy categories, the temperature-dependent drops in strength are also composition/microstructure dependent. The reasons for such behaviors are currently not identified. The purpose of the present work is to evaluate the influence of alloy melting temperature and the volume fraction of second phases on the temperature dependence of RCCA yield strength. To explore these effects, we analyze the behavior of 15 single-phase and 11 multi-phase RCCAs for which the mechanical properties at room temperature (RT), 1000°C and 1200°C have been reported [1, 6]. Using the CALPHAD approach, we calculate phase diagrams for these same alloys to identify the melting (T_m) temperatures, volume fractions of secondary phases and the solvus temperatures (T_s) at which the secondary phases dissolve in the multi-phase RCCAs studied here. We then explore correlations between the strength drops at 1000°C and 1200°C, the alloy compositions, melting temperatures and volume fractions of secondary phases and analyze the results. We also explore the influence of alloy density on the temperature dependence of specific yield strength. Finally, based on this analysis we make several conclusions, which may be useful for the future development of high-temperature RCCAs.

2. METHODS

15 single-phase RCCAs and 11 multi-phase RCCAs, for which compression properties at RT, 1000°C and 1200°C have been reported in the open literature or acquired by the authors, have been selected for this study. These alloys are drawn from a palette of 12 elements (Al, Cr, Hf, Mo, Nb, Re, Si, Ta, Ti, V, W, and Zr). They were prepared by mixing and arc-melting high-purity elements followed by solidification on a copper hearth. The compositions of these alloys and their yield strength values are given in Table 1. Nine alloys were in the as-solidified (AC) condition and seventeen alloys were in the annealed (A) condition. For fourteen alloys, hot isostatic pressing (HIP) preceded the annealing step. In spite of different processing conditions, all the studied alloys had coarse-grained structure, with the average grain size exceeding 100 μm . At such large grain sizes the grain boundary effects on the yield strength and its temperature dependence are considered to be negligible in comparison with solid-solution and precipitation

strengthening [8-10]. Therefore, grain boundary (Hall-Petch) strengthening is not discussed in this work.

Table 1. Composition, processing condition, phase content, density and yield strength, determined at 23°C, 1000°C and 1200°C, of selected RCCAs. Data are taken from [1] and from the present work.

Alloy	Condition [#]	Phases	Density g/cm ³	YS at 23°C (MPa)	YS at 1000°C (MPa)	YS at 1200°C (MPa)
Reported single-phase alloys						
Al _{0.4} Hf _{0.6} NbTaTiZr	HIP+A	BCC	9.08	1841	298	89
AlCrMoTi	A	BCC	5.98	1100	375	100
AlMoNbTi	A	BCC	6.46	1100	540	200
HfMo _{0.5} NbTiV _{0.5}	AC	BCC	8.96	1260	368	60
HfMoNbTaTiZr	AC	BCC	9.93	1512	814	556
HfMoNbTiZr	AC	BCC	8.68	1575	635	187
HfMoTaTiZr	AC	BCC	10.18	1600	855	404
HfNbTaTiZr	HIP+A	BCC	9.89	929	295	92
MoNbTaTiVW	AC	BCC	10.91	1515	753	659
MoNbTaTiW	AC	BCC	11.72	1343	620	586
MoNbTaVW	HIP+A	BCC	12.29	1246	842	735
MoNbTaW	HIP+A	BCC	13.6	996	548	506
MoNbTi	HIP+A	BCC	7.67	1100	504	324
NbTa _{0.3} TiZr	HIP+A	BCC	7.81	882	274	102
NbTiZr	HIP+A	BCC	6.63	975	141	61
Reported multi-phase alloys						
AlCrMoNbTi	A	BCC+Laves	6.56	1010	550	105
Al _{0.24} NbTiVZr	HIP+A	BCC+Laves	6.34	1240	82	24
AlMo _{0.5} NbTa _{0.5} TiZr	HIP+A	B2+BCC+Al ₄ Zr ₅	7.14	2000	745	250
Cr _{0.3} NbTiZr	HIP+A	BCC+Laves	6.56	1576	139	32
CrMo _{0.5} NbTa _{0.5} TiZr	HIP+A	BCC+BCC+Laves	7.98	1595	546	170
HfMo _{0.5} NbSi _{0.3} TiV _{0.5}	AC	BCC+M ₅ Si ₃	8.46	1617	398	166
HfMo _{0.5} NbSi _{0.5} TiV _{0.5}	AC	BCC+M ₅ Si ₃	8.16	1787	614	188
HfMo _{0.5} NbSi _{0.7} TiV _{0.5}	AC	BCC+M ₅ Si ₃	7.89	2134	673	235
MoNbTiVZr	HIP+A	BCC+Laves	7.27	1706	467	116
NbRe _{0.3} TiZr	HIP+A	BCC+Laves	7.81	1244	323	89
NbTaTiVZr	HIP+A	BCC+BCC	8.43	1395	190	78

Hot isostatic pressing (HIP), Annealed (A), As-cast (AC)

Phase diagrams for the studied alloys were calculated using the CALPHAD approach. We used ThermoCalc software and the TCHEA3-2018 thermodynamic database developed by ThermoCalc [11, 12] and Pandat software (version 2018) and the PanNb2018a database

developed by CompuTherm, LLC [13, 14] for the calculations. The TCHEA3-2018 database contains 26 elements (Al, B, C, Co, Cr, Cu, Fe, Hf, Ir, Mn, Mo, N, Nb, Ni, Re, Rh, Ru, Si, Sn, Ta, Ti, V, W, Y, Zn, and Zr). All binary systems and 25 ternary systems from the 12-elements included in this study are assessed to their full range of composition in this database. The PanNb2018a database contains 13 elements (Al, Cr, Fe, Hf, Mo, Nb, Re, Si, Ta, Ti, V, W, and Zr) and 135 phases. It has complete thermodynamic descriptions for all the binary systems and 34 ternary systems, among which 21 are related to the alloy systems explored in this work.

3. RESULTS

3.1 Calculation of Phase Diagrams of the Studied Alloys

The level of correctness of the phase diagram calculations and the respective thermodynamic properties directly depend on the level of development of the used thermodynamic database. The calculation for a given alloy is considered to be the most reliable if the database has complete thermodynamic descriptions for all binary and ternary systems included in that alloy [15]. Both the PanNb2018a and TCHEA3-2018 have complete descriptions of the binary systems, but only few of the ternary systems related to the studied alloys are assessed. When the ternary systems are unavailable, thermodynamic calculations are extrapolated from the binary systems [16-18], and such calculations should be treated with caution [17-19]. The credibility of CALPHAD calculations increases with an increase in the fractions of the fully assessed ternary (FAT) systems. FAT is calculated as the ratio of the number of fully assessed ternaries for a given alloy system to the total number of ternaries in that alloy system [15]. FAT values for each of the studied alloys in the PanNb2018a and TCHEA3-2018 databases are given in Table 2. The PanNb2018a database has higher FAT values for 21 of the 26 studied alloys, while TCHEA3-2018 has higher credibility for only two alloys, AlCrMoTi and AlCrMoNbTi. Both databases have the same credibility for 3 alloys, AlMoNbTi, CrMo_{0.5}NbTa_{0.5}TiZr and MoNbTaW. Nevertheless, for completeness, both databases were used to calculate phase diagrams for all the studied alloys and the results are compared with the available experimental data. The calculated phase diagrams are given in Supplementary Materials. The calculated solidus (T_m) and solvus (T_s) temperatures, as well as solid phases at T_m and a T_s -phase, are given in Table 2. T_s is identified as the highest phase transition temperature before melting, and the T_s -phase is a phase which dissolves at T_s upon heating.

Table 2. Fractions of the assessed ternary systems (FAT) in PanNb2018a (Pan) and TCHEA3-2018 (TC) databases for the studied alloys. Also shown for both databases are the solidus (T_m) and solvus (T_s) temperatures; the phases present at T_m ; the phase related to T_s ; and T_m calculated using the rule of mixtures (ROM). Preferred calculated values are shown in bold font (see Sec. 3.1.1).

Alloy	FAT Pan	FAT TC	T_m , °C Pan	T_m , °C TC	T_m , °C ROM	Phases at T_m Pan	Phases at T_m TC	T_s , °C Pan	T_s , °C TC	T_s -phase Pan	T_s -phase TC
Reported single-phase alloys											
Al _{0.4} Hf _{0.6} NbTaTiZr	0.3	0.15	1732	1632	2124	BCC1	BCC1	1636	1356	BCC2	BCC2
AlCrMoTi	0	1	1663	1876	1715	BCC1	BCC	975	931	A15	A15
AlMoNbTi	0.5	0.5	1959	1780	1857	BCC1	BCC	846	1143	A15	A15
HfMo _{0.5} NbTiV _{0.5}	0.2	0	1987	1776	2161	BCC1	BCC1	724	827	BCC2	BCC2
HfMoNbTaTiZr	0.3	0	1860	1840	2312	BCC1	BCC1	1507	1171	BCC2	BCC2
HfMoNbTiZr	0.3	0	1844	1767	2171	BCC1	BCC1	917	1038	BCC2	BCC2
HfMoTaTiZr	0.1	0	1723	1793	2279	BCC1+BCC2	BCC1	906	1168	HCP	BCC2
HfNbTaTiZr	0.5	0	1882	1851	2250	BCC1	BCC1	984	1019	BCC2	BCC2
MoNbTaTiVW	0.2	0	2182	2296	2520	BCC1	BCC	846	572	BCC2	HCP
MoNbTaTiW	0.3	0	2395	2416	2641	BCC1	BCC1	1100	708	BCC2	BCC2
MoNbTaVW	0.1	0	2409	2566	2690	BCC1	BCC1	824	410	BCC2	BCC2
MoNbTaW	0	0	2807	2804	2885	BCC	BCC	*	*	*	*
MoNbTi	1	0	2115	1970	2256	BCC1	BCC1	740	781	BCC2	BCC2
NbTa _{0.3} TiZr	0.75	0	1711	1727	2092	BCC1	BCC1	850	831	BCC2	BCC2
NbTiZr	1	0	1688	1691	2000	BCC1	BCC1	600	694	BCC2	BCC2
Reported multi-phase alloys											
AlCrMoNbTi	0.5	0.8	1721	1780	1867	BCC1	BCC1	840	1534	C14	C14
Al _{0.24} NbTiVZr	0.2	0.1	1545	1491	1903	BCC1	BCC1	887	863	BCC2	BCC2
AlMo _{0.5} NbTa _{0.5} TiZr	0.3	0.25	1595	1478	1896	BCC1+BCC2	BCC1+BCC2+σ	1197	1200	Al₄Zr₅	Al ₄ Zr ₅
Cr _{0.3} NbTiZr	1	0.5	1465	1410	1991	BCC1	BCC	1185	1273	C15	C15
CrMo _{0.5} NbTa _{0.5} TiZr	0.45	0.45	1397	1400	2145	BCC1+C15	BCC1+C15	1347	1050	BCC2	BCC2
HfMo _{0.5} NbSi _{0.3} TiV _{0.5}	0.35	0.2	1621	1190	2109	BCC1+M₅Si₃	BCC1+C16	1114	750	M₂Si	BCC2
HfMo _{0.5} NbSi _{0.5} TiV _{0.5}	0.35	0.2	1630	1213	2078	BCC1+ M₅Si₃	BCC1+C16+ M ₅ Si ₃	1118	690	M₂Si	M ₃ Si
HfMo _{0.5} NbSi _{0.7} TiV _{0.5}	0.35	0.2	1642	1222	2050	BCC1+ M₅Si₃	BCC1+C16+ M ₅ Si ₃	1120	620	M₂Si	M ₃ Si
MoNbTiVZr	0.2	0	1649	1583	2107	BCC1	BCC1	994	970	BCC2	BCC2
NbRe0.3TiZr	0.75	0.5	1704	1683	2108	BCC1	BCC	1656	933	C14	ReZr ₂
NbTaTiVZr	0.4	0	1612	1581	2185	BCC1	BCC1	1036	942	BCC2	BCC2

* No secondary phase is predicted to form for this alloy

Analysis of the calculations show that, for the same alloy, the two databases calculate almost the same T_m values, with the exception of three Si-containing alloys, for which T_m^{TC} values were considerably lower than T_m^{Pan} (Figure 1a). The superscripts ‘TC’ and ‘Pan’ indicate that the

quantities were calculated using ThermoCalc or Pandat software, respectively. It is likely that the TC calculations for the Si-containing alloys are incorrect, because they predict melting of these alloys around 1200°C, while the experimental results revealed that these alloys were solid and showed rather high strength at 1200°C. With five exceptions, the Pan and TC calculations predict almost the same phases at T_m , for each of the studied alloys (Table 2). However, the T_s values calculated with PanNb2018a and TCHEA3-2018 are quite different (Figure 1b). In spite of this, both databases calculate the same T_s -phases for each of the studied alloys, with only three exceptions (Table 2).

Figure 1. Comparison of (a) solidus (T_m^{Pan} , T_m^{TC}) and (b) solvus (T_s^{Pan} , T_s^{TC}) temperatures calculated with PanNb2018a (Pan) and TCHEA3-2018 (TC) thermodynamic databases.

3.1.1 Comparison with experiment and CALPHAD calculations selection

Single-phase RCCAs

For the 15 RCCAs that are experimentally reported as single-phase BCC structures, both PanNb2018a and TCHEA3-2018 predict single-phase BCC phase field with a large temperature interval between T_m and T_s (Table 2). Formation of equilibrium secondary phases is predicted at temperatures below 1000°C in all but one of these alloys. Therefore, the single phase BCC structures observed at room temperature are likely metastable because formation of secondary phases that are thermodynamically stable at room temperature is kinetically restricted by low T_s and slow diffusion. These supersaturated solid solutions may decompose and form secondary phases when heated for sufficient time between 23°C and T_s . Only one alloy, MoNbTaW, is

predicted by both databases to be a thermodynamically stable single-phase BCC structure at room temperature.

Comparison of the CALPHAD calculations with experimental results allows identification of a preferred database for each of the studied alloys. First, for $\text{Al}_{0.4}\text{Hf}_{0.6}\text{NbTaTiZr}$ and HfMoNbTaTiZr PanNb2018a predicts formation of a second phase (BCC2) at relatively high temperatures, $T_s = 1636^\circ\text{C}$ and 1507°C , respectively. If the calculations are correct, the alloys should be two-phase structures at room temperature, which contradict experiments. On the other hand, TCHEA3-2018 predicts much lower $T_s = 1356^\circ\text{C}$ and 1171°C , respectively, for these alloys. Experiments do not show a second phase in the 1200°C annealed $\text{Al}_{0.4}\text{Hf}_{0.6}\text{NbTaTiZr}$ [20] and as-cast HfMoNbTaTiZr [21], indirectly indicating that T_s is low enough to restrict the phase transformation kinetically. As a result, TCHEA3-2018 data will be used for these two alloys. Second, PanNb2018a predicts two BCC phases in HfMoTaTiZr after solidification, which disagrees with the experiment. On the other hand, TCHEA3-2018 predicts one BCC phase at T_m and the second BCC phase forms in equilibrium conditions below $T_s = 1168^\circ\text{C}$. Therefore, the results from TCHEA3-2018 calculations will be used for HfMoTaTiZr . Finally, MoNbTaTiW has been found experimentally to be a single-phase BCC structure both in the as-cast condition and after annealing at 1200°C for 24 h [22], which agrees better with TCHEA3-2018 than with PanNb2018a calculations (Table 2).

Multi-phase RCCAs

Eight RCCAs that do not contain Si are reported to be multi-phase after annealing at 1200°C or 1400°C , and 3 Si-containing RCCAs are reported as multi-phase in the as-cast condition (Table 1). CALPHAD predictions of the type and number of phases in these alloys (Table 2) agree well with the experimental data. Both PanNb2018a and TCHEA3-2018 databases predict that 5 of these multi-phase RCCAs ($\text{AlMo}_{0.5}\text{NbTa}_{0.5}\text{TiZr}$, $\text{CrMo}_{0.5}\text{NbTa}_{0.5}\text{TiZr}$, $\text{HfMo}_{0.5}\text{NbSi}_{0.3}\text{TiV}_{0.5}$, $\text{HfMo}_{0.5}\text{NbSi}_{0.5}\text{TiV}_{0.5}$ and $\text{HfMo}_{0.5}\text{NbSi}_{0.7}\text{TiV}_{0.5}$) are not single-phase above T_s and retain a multi-phase structure up to T_m . The other 6 RCCAs have a single-phase BCC region above T_s ; e.g. $\text{Al}_{0.24}\text{NbTiZr}$, MoNbTiVZr and NbTaTiVZr have T_s below 1200°C , according to both databases, and thus they can be considered as single-phase structures when tested above T_s .

Selection of CALPHAD calculations for mechanical property analysis

In the analyses conducted in the following sections, CALPHAD calculations that best agree with the reported experimental data are used. When both PanNb2018a and TCHEA3-2018 calculations agree with the experimental results equally well, the calculations with higher FAT values are used. If FAT values are the same, TCHEA3-2018 calculations are used for the alloys which do not contain Nb. The T_s and T_m values and respective phases thus selected are highlighted by a bold font in Table 2 for each of the studied alloys. The superscript ‘CALPHAD’ or no superscript is used to indicate these preferred values in the remainder of this work.

3.1.2 Comparison of CALPHAD-calculated and rule-of-mixtures calculated T_m

Figure 2 compares T_m^{CALPHAD} values for the studied alloys with those calculated using rule of mixtures (ROM):

$$T_m^{\text{ROM}} = \sum c_i T_{mi} \quad (1)$$

Figure 2. Comparison of the CALPHAD-calculated solidus temperatures T_m^{CALPHAD} for the studied alloys with those calculated with the use of the rule of mixtures, T_m^{ROM} . The dashed line corresponds to the condition when $T_m^{\text{ROM}} = T_m^{\text{CALPHAD}}$.

Here c_i is the atomic fraction and T_{mi} is the melting temperature of element i . It is found that T_m^{ROM} is higher than T_m^{CALPHAD} for almost all the alloys. The difference is generally larger for alloys with lower T_m , i.e. for alloys containing Al and Cr, as well as for multi-phase alloys (Table 2). Thus, the ROM overestimates T_m for the studied RCCAs. Such behavior can be expected

because ROM predicts a straight solidus line between any two melting points, while in the solid-solution based alloy systems, with no high-temperature compounds, the solidus line always deviates from a straight line towards lower temperatures.

3.2 Effect of the melting temperature on the strength of RCCAs

It can be seen from the CALPHAD calculations (Table 2) that the multi-phase RCCAs have relatively low T_m , often considerably lower than single-phase RCCAs. Indeed, T_m of the reported multi-phase RCCAs varies from 1397°C for CrMo_{0.5}NbTa_{0.5}TiZr to 1780°C for AlCrMoNbTi, while T_m of the reported single-phase RCCAs varies from 1632°C for Al_{0.4}Hf_{0.6}NbTaTiZr to 2804°C for MoNbTaW. Only two single-phase alloys have T_m below 1700°C, while only two multi-phase alloys have T_m above 1700°C. The lower melting temperatures can limit operating temperatures of the multi-phase RCCAs relative the single-phase RCCAs. The behavior of these two categories of RCCAs at different temperatures is described in detail in the following sections.

Room temperature (RT) behavior

There is no clear correlation between T_m and the yield strength of RCCAs at 25°C (Figure 3a). Single-phase alloys have RT yield strength $\sigma_{0.2}^{25^\circ\text{C}}$ in the range from 882 MPa for NbTa_{0.3}TiZr ($T_m = 1711^\circ\text{C}$) to 1841 MPa for Al_{0.4}Hf_{0.6}NbTaTiZr ($T_m = 1632^\circ\text{C}$). The RT yield strength of MoNbTaW, which has the highest T_m ($= 2804^\circ\text{C}$), is 996 MPa. The multi-phase alloys have, on average, higher RT yield strength than the single-phase alloys (Figure 3a). The $\sigma_{0.2}^{25^\circ\text{C}}$ of these alloys is in the range from 1010 MPa for AlCrMoNbTi ($T_m = 1780^\circ\text{C}$) to 2134 MPa for HfMo_{0.5}NbSi_{0.7}TiV_{0.5} ($T_m = 1642^\circ\text{C}$). Figure 3b shows the dependence of $\sigma_{0.2}^{25^\circ\text{C}}$ on the atomic radius misfit parameter, δ_r . By definition, $\delta_r = 100\% \sqrt{\sum c_i (1 - \frac{r_i}{\bar{r}})^2}$, where c_i and r_i are the mole fraction and atomic radius of an alloying element i , respectively, and $\bar{r} = \sum c_i r_i$. It can be seen that the single-phase RCCAs have $\delta_r < 7.5\%$ and there is no clear correlation between $\sigma_{0.2}^{25^\circ\text{C}}$ and δ_r for these alloys. The multi-phase RCCAs have δ_r from 5.3 to 10.2% and there is a tendency for the alloys containing the Laves and silicide phases to increase $\sigma_{0.2}^{25^\circ\text{C}}$ with increasing δ_r . Figure 3c shows the dependence of $\sigma_{0.2}^{25^\circ\text{C}}$ on the experimentally reported volume fraction of all of the secondary phases, V , in multi-phase alloys. There is a clear tendency for

$\sigma_{0.2}^{25^\circ\text{C}}$ to increase with an increase in the volume fraction of the secondary phases; however, the alloy chemical composition (i.e. atom interaction forces) is also important. For example, $\text{Al}_{0.24}\text{NbTiVZr}$ and MoNbTiVZr have almost the same volume fraction (36% and 35%, respectively) of the secondary Laves phase; however $\sigma_{0.2}^{25^\circ\text{C}}$ of the first alloy is 1240 MPa and the second alloy is 1706 MPa. This analysis indicates that the higher RT strengths of the multi-phase alloys are likely a result of additional strengthening from the multi-phase microstructures and increasing volume fraction of the secondary phases.

Figure 3. Room temperature yield strength ($\sigma_{0.2}^{25^\circ\text{C}}$) of single-phase (S) and multi-phase (M) RCCAs as a function of (a) the alloy melting temperature T_m , (b) atomic radius misfit parameter δ_r , and (c) volume fraction (in percent) of the secondary phases. The type of the major secondary phase is shown in the legends.

Behavior of RCCAs at 1000°C

According to CALPHAD calculations and their comparison with the experimental data, the single-phase BCC RCCAs are metastable supersaturated solid solutions at room temperature. Therefore, during heating and holding at a testing temperature (1000°C) they may experience phase transformations and become multi-phase structures. The diffusion processes required to initiate the phase transformations generally become noticeable at temperatures $T > 0.4T_m$, where T_m is the absolute melting temperature, while at $T < 0.4T_m$ the phase transformations are kinetically restricted [23]. It is therefore assumed in this work that the single-phase RCCAs with $T_s > 1000^\circ\text{C}$ and $T_m < 2910^\circ\text{C}$ (3183 K) become multi-phase structures during heating and deformation at 1000°C. Four RCCAs, $\text{Al}_{0.4}\text{Hf}_{0.6}\text{NbTaTiZr}$, AlMoNbTi , HfMoNbTaTiZr and HfMoTaTiZr , satisfy these conditions. Other single-phase RCCAs retain their single-phase structure at 1000°C. For the same reason, RCCAs that are multi-phase at 25°C and also have $T_s \leq 1000^\circ\text{C}$, $T_m < 2910^\circ\text{C}$ (3183 K) and a single phase at T_m , become single-phase structures at 1000°C. These are $\text{Al}_{0.24}\text{NbTiVZr}$ and MoNbTiVZr . Other multi-phase RCCAs retain their multi-phase structure at 1000°C.

Figure 4a shows the dependence of the yield strength at 1000°C ($\sigma_{0.2}^{1000^\circ\text{C}}$) on T_m for single-phase and multi-phase RCCAs. There is clear tendency for single-phase RCCAs to increase in strength at 1000°C with increasing T_m . For example, alloys with $T_m < 1750^\circ\text{C}$, between 1750°C and 2000°C and above 2000°C have $\sigma_{0.2}^{1000^\circ\text{C}}$ in the range from 141 MPa to 274 MPa, from 295 MPa to 635 MPa, and from 504 MPa to 842 MPa, respectively. The four RCCAs observed to be single-phase at 25°C but predicted by CALPHAD to become multi-phase at 1000°C show a trend to increase $\sigma_{0.2}^{1000^\circ\text{C}}$ beyond the typical range for single-phase RCCAs with the same T_m values due to additional precipitation-strengthening effect.

The multi-phase RCCAs show no dependence of $\sigma_{0.2}^{1000^\circ\text{C}}$ on T_m . These alloys have a narrow T_m range and the alloys with almost the same T_m can have quite different $\sigma_{0.2}^{1000^\circ\text{C}}$ values ranging from 139 MPa for $\text{Cr}_{0.3}\text{NbTiZr}$ to 745 MPa for $\text{AlMo}_{0.5}\text{NbTa}_{0.5}\text{TiZr}$ (Figure 4a, Table 1). A former multi-phase $\text{Al}_{0.24}\text{NbTiVZr}$, which contains a Laves phase at room temperature and has $\sigma_{0.2}^{25^\circ\text{C}} = 1240$ MPa, becomes a single-phase BCC structure with the lowest strength ($\sigma_{0.2}^{1000^\circ\text{C}} = 82$ MPa) at 1000°C among all the reported RCCAs, probably due to its relatively low $T_m = 1545^\circ\text{C}$ (Figure 4a). On the other hand, MoNbTiVZr also becomes single-phase at 1000°C but it has a T_m of $\sim 100^\circ\text{C}$ higher than the former alloy and retains a relatively high strength of $\sigma_{0.2}^{1000^\circ\text{C}}$

= 467 MPa at 1000°C. The behaviors of these two alloys are in line with the behavior of single-phase RCCAs (Figure 4a).

Figure 4. (a) Yield strength at 1000°C ($\sigma_{0.2}^{1000^\circ\text{C}}$) and (b) yield strength ratio ($\sigma_{0.2}^{1000^\circ\text{C}}/\sigma_{0.2}^{25^\circ\text{C}}$) of single-phase (S) and multi-phase (M) RCCAs as a function of the melting temperature T_m .

Figure 4b compares yield strength values at 1000°C and 25°C of the single-phase and multi-phase RCCAs with different melting temperatures. A strength drop occurs between these temperatures and the magnitude of the drop, $\eta_1 = \sigma_{0.2}^{1000^\circ\text{C}}/\sigma_{0.2}^{25^\circ\text{C}}$, strongly depends on the type of alloy. For the single-phase RCCAs, η_1 is the lowest (=0.07) for the $\text{Al}_{0.24}\text{NbTiVZr}$, which has the lowest $T_m = 1545^\circ\text{C}$. The ratio tends to increase with increasing T_m and approaches $\eta_1 = 0.45$ -0.7 for single-phase alloys with $T_m > 2000^\circ\text{C}$. On the other hand, the η_1 values for multi-phase RCCAs do not depend on T_m and for alloys with almost the same T_m η_1 varies from 0.09 to 0.54. Figure 5 shows dependence of $\sigma_{0.2}^{1000^\circ\text{C}}$ of the multi-phase alloys on (a) the volume fraction of the secondary phases V at 1000°C and (b) the temperature difference between the solvus, T_s , and testing temperatures ($\Delta T = T_s - 1000^\circ\text{C}$). It can be clearly seen from Figure 5a that the different strength values of multi-phase alloys at 1000°C are due to the different volume fractions of the secondary phases. With the same type of secondary phase, $\sigma_{0.2}^{1000^\circ\text{C}}$ increases with an increase in V . Because V generally decreases with increasing temperature and approaches zero at $T = T_s$, $\sigma_{0.2}^{1000^\circ\text{C}}$ tends to increase with increasing T_s (ΔT), Figure 5b. Because different alloys have different dependence of V on ΔT , the relationship between $\sigma_{0.2}^{1000^\circ\text{C}}$ and ΔT is not as straightforward as that between $\sigma_{0.2}^{1000^\circ\text{C}}$ and V , when different multi-phase alloys are

considered, and the alloys with similar ΔT may have quite different $\sigma_{0.2}^{1000^\circ\text{C}}$ values (compare Figure 5a and Figure 5b).

These results clearly show that the drop in strength between 25°C and 1000°C is more dramatic for single-phase RCCAs with lower T_m and for multi-phase RCCAs with a stronger temperature dependence of the volume fraction of secondary phases. Single-phase RCCAs with $T_m > 2000^\circ\text{C}$, or multi-phase RCCAs with $T_s > 1100^\circ\text{C}$ and the secondary-phase volume fraction $\geq 20\%$ at 1000°C , have better ability to retain high strength at 1000°C .

Figure 5. The yield strength ($\sigma_{0.2}^{1000^\circ\text{C}}$) of the multi-phase (M) RCCAs with different secondary phases as a function of (a) the volume fraction V of the secondary phases at 1000°C and (b) the temperature difference between the solvus (T_s) and testing (1000°C) temperatures.

Behavior of RCCAs at 1200°C

All but one of the 15 RCCAs that are reported to be single-phase BCC at 25°C have $T_s < 1200^\circ\text{C}$ and are therefore thermodynamically stable single-phase structures at 1200°C . The exception is $\text{Al}_{0.4}\text{Hf}_{0.5}\text{NbTaTiZr}$, which, in accord to CALPHAD calculations, has a BCC1+BCC2 structure at 1200°C . Among the 11 RCCAs reported to be multi-phase at 25°C , $\text{Al}_{0.24}\text{NbTiVZr}$, $\text{Cr}_{0.3}\text{NbTiZr}$, MoNbTiVZr and NbTaTiVZr become single-phase BCC structures at 1200°C and the other 7 alloys retain their multi-phase structure. The dependence of the yield strength at 1200°C ($\sigma_{0.2}^{1200^\circ\text{C}}$) on T_m for the single-phase and multi-phase RCCAs is shown in Figure 6a. There is a general trend for $\sigma_{0.2}^{1200^\circ\text{C}}$ of the RCCAs that are single-phase structures at 1200°C to increase with increasing T_m ; however 3 alloys with moderate T_m values, AlCrMoTi ($T_m = 1876^\circ\text{C}$), HfNbTaTiZr ($T_m = 1882^\circ\text{C}$) and $\text{HfMo}_{0.5}\text{NbTiV}_{0.5}$ ($T_m = 1987^\circ\text{C}$), have $\sigma_{0.2}^{1200^\circ\text{C}}$ less than 100 MPa, which is typical for alloys with $T_m < 1750^\circ\text{C}$. The overall trend of $\sigma_{0.2}^{1200^\circ\text{C}}$ with

T_m can be described by a band:

$$\sigma_{0.2}^{1200^\circ\text{C}} = 0.625(T_m - T_B) \quad (2)$$

where T_B is in the range from 1100°C to 1700°C (Figure 6a).

The $\sigma_{0.2}^{1200^\circ\text{C}}$ values of the multi-phase RCCAs are similar to those of the single-phase RCCAs with the same (low) T_m values (Figure 6a). Therefore, multi-phase RCCAs have considerably lower strength than many single-phase RCCAs with higher T_m .

Softening behavior in the temperature range between 1000°C and 1200°C of the studied RCCAs with different T_m values is shown in Figure 6b. There are two groups of single-phase RCCAs with different dependence of the strength drop $\eta_2 = \sigma_{0.2}^{1200^\circ\text{C}}/\sigma_{0.2}^{1000^\circ\text{C}}$ on T_m . In the first group, which includes AlMoNbTi, HfNbTaTiZr, HfMoNbTiZr, AlCrMoTi and HfMo_{0.5}NbTiV_{0.5}, η_2 decreases (from 0.372 to 0.163) with increasing T_m (from 1711°C to 1987°C). In the second group, which includes all other single-phase RCCAs, η_2 increases with increasing T_m . The ‘anomalous’ behavior of the stress ratio η_2 for the first group of the single-phase RCCAs can be explained if the presence of non-equilibrium secondary phases (BCC2 in the three Hf-containing and A15 in two Al-containing RCCAs), which precipitate during heating and holding at 1200°C, is assumed. Although T_s for these alloys is smaller than the testing temperature (1200°C), the secondary phases may remain at 1200°C due to insufficient diffusion kinetics and cause additional strengthening. In this case, a higher volume fraction of a residual secondary phase is expected for an alloy with smaller $\Delta T = (1200^\circ\text{C} - T_s)$. Indeed, $\sigma_{0.2}^{1200^\circ\text{C}}/\sigma_{0.2}^{1000^\circ\text{C}}$ increases with decreasing ΔT for these 5 RCCAs (Figure 6c).

The multi-phase RCCAs do not show any dependence of $\sigma_{0.2}^{1200^\circ\text{C}}/\sigma_{0.2}^{1000^\circ\text{C}}$ on T_m (Figure 6b). However, their strength drop relates directly to the decrease in the volume fraction of the secondary phases (Figure 6d).

Figure 6. (a) Yield strength at 1200°C ($\sigma_{0.2}^{1200^\circ\text{C}}$) and (b-d) yield strength ratio ($\sigma_{0.2}^{1200^\circ\text{C}} / \sigma_{0.2}^{1000^\circ\text{C}}$) of single-phase (S) and multi-phase (M) RCCAs as a function of (b) the melting temperature T_m or (c) the difference between testing (1473 K) and equilibrium solvus (T_s) temperatures (for five single-phase alloys showing ‘anomalous’ dependence on T_m) normalized by T_m or (d) the volume fraction ratio ($V^{1200^\circ\text{C}} / V^{1000^\circ\text{C}}$) of the secondary phases in multi-phase RCCAs. The bubble in (a) represents the three off-trend alloys, i.e. HfNbTaTiZr, AlCrMoTi and HfMo_{0.5}NbTiV_{0.5}. The bubble in (b) shows the five reverse-trend alloys, i.e. AlMoNbTi and HfMoNbTiZr in addition to the three previous ones in (a).

4. DISCUSSION

4.1 Influence of the T_m on the Temperature Dependence of RCCA Yield Strength

Both the CALPHAD calculations and ROM analysis show that the reported single-phase RCCAs have a wide spread in T_m ranging, according to CALPHAD, from 1632°C for Al_{0.4}Hf_{0.6}NbTaTiZr to 2807°C for MoNbTaW. The T_m range of the reported multi-phase RCCAs is much narrower, from 1397°C for CrMo_{0.5}NbTa_{0.5}TiZr to 1780°C for AlCrMoNbTi. Formation of secondary

phases during solidification is associated with considerably lower T_m values.

Single-phase RCCAs show RT yield strength values, $\sigma_{0.2}^{25^\circ\text{C}}$, in the range from 880 MPa to 1790 MPa. No clear relationship is found between $\sigma_{0.2}^{25^\circ\text{C}}$ and T_m or δ_r for the reported RCCAs. This agrees with recent work by Rao *et al.* [24, 25] who used atomistic simulations and the Suzuki model to calculate the yield strengths of 3 ternary and 9 quaternary BCC RCCAs. They showed that the solid solution strengthening of these alloys at temperatures where diffusion processes are not active, including room temperature, is likely controlled by the stacking fault energy fluctuations along screw dislocations rather than by the atomic radius or shear modulus misfits. These stacking fault energy fluctuations cause significant barriers for kink migration and for dragging edge dislocation dipoles produced by kink collisions along the screw dislocations. The solute-dislocation interaction energies are composition dependent, which explains large variations in the RT yield strengths for the single-phase RCCAs (Figure 3a). In addition to solid-solution strengthening, multi-phase RCCAs are also strengthened by secondary phases. As a result, $\sigma_{0.2}^{25^\circ\text{C}}$ of multi-phase RCCAs tends to increase with increasing V , and alloys with $V \geq 40\%$ are generally stronger than single-phase RCCAs.

Other strengthening mechanisms, such as grain boundary (Hall-Petch) or dislocation forest strengthening provide second-order effects on yield strength. All the studied alloys have coarse-grained structures, with the average grain size exceeding 100 μm . Taking a typical value of 400 $\text{MPa } \mu\text{m}^{-0.5}$ for the Hall-Petch constant k_σ [8], the contribution of the grain size d to the yield strength is ~ 40 MPa or 28 MPa when $d = 100$ μm or 200 μm , which is negligible compared to the reported yield strength values. Recent analysis of the Hall-Petch effect on a classical refractory HEA HfNbTaTiZr resulted in $k_\sigma = 270$ $\text{MPa } \mu\text{m}^{-0.5}$ [10]. This value of the Hall-Petch constant gives even smaller strengthening contribution from grain boundaries, only 27 MPa at $d = 100$ μm . The contribution to the yield strength from forest dislocations is also low (< 50 MPa) as the dislocation density in HIP+A, A or AC conditions is low ($\leq 10^{12} \text{m}^{-2}$). Strong texture is generally formed after extensive deformation and it is not typical in cast and annealed alloys. Microsegregation may affect local microhardness, but the macroscopic strength is generally unaffected. Thus, the strengths of the reported RCCAs are largely controlled by dislocation/solute and dislocation/precipitate interactions.

The solid solution strengthening model by Rao *et al.* [24, 25] does not take into account diffusion effects and it agrees well with the experimental data only at relatively low temperatures, where

diffusion-controlling deformation mechanisms can be ignored. Earlier analysis of the deformation behavior of a HfNbTaTiZr RCCA showed at least two strengthening mechanisms: (i) thermally-activated, diffusionless dislocation glide (similar to that described above) at low temperatures and (ii) diffusion-controlled dislocation climb/glide at high temperatures [26]. The diffusion-controlled regime generally occurs above $\sim 0.5\text{--}0.6T_m$ and can be distinguished from the low-temperature regime by a more rapid drop in strength with increasing temperature, because dislocations are able to move more easily around obstacles [27, 28]. To identify the location of this transition in the studied RCCAs, their $\log(\sigma_{0.2})$ values are plotted versus T/T_m in Figure 7a and Figure 7b for the single-phase and multi-phase RCCAs, respectively. The strength data for additional temperatures, taken from [6] or obtained by the authors, are also shown for several alloys to better identify the transition. For comparison, curves are shown in Figure 7a for commercial alloys based on Mo (TZM) and Nb (C-103), and for pure tungsten.

Two temperature regions, with very different temperature dependence of the yield strength, are clearly seen. The transition from the low-temperature regime to the high-temperature regime is manifested by a stronger decrease in the yield strength with increasing temperature. All the single-phase RCCAs show the transition at almost the same homologous temperature of $\sim 0.6 T_m$ (Figure 7a). In seven single-phase RCCAs, which have $T_m < 2121$ K (1848°C), this transition occurs below 1000°C. In four single-phase RCCAs, with T_m between 2143 and 2393 K (1870 and 2120°C), it occurs between 1000°C and 1200°C. Only four remaining single-phase RCCAs (MoNbTaW, MoNbTaTiW, MoNbTaVW and MoNbTaTiVW) for which $T_m \geq 2455$ K (2182°C) show transition to the diffusion-controlling regime at temperatures $\geq 1200^\circ\text{C}$. Since $\sigma_{0.2}$ drops more quickly with increasing T/T_m in the diffusion-controlled regime than in the glide-controlling regime, this explains why the single-phase RCCAs with higher T_m are generally stronger at 1000°C and 1200°C. For comparison, the transition to the high-temperature regime in pure tungsten occurs at $\sim 0.45 T_m$ (1371°C) and in TZM and C-103 at $\sim 0.55 T_m$ (1300°C and 1090°C, respectively) and these conventional materials have noticeably lower strength than many studied RCCAs at equivalent T/T_m (Figure 7a). A clear advantage of the RCCAs relative to the conventional refractory alloys can be seen: RCCAs are stronger and they have higher high-temperature capability.

The strength of multi-phase RCCAs is mainly controlled by solid-solution, precipitation and dispersoid strengthening, with contributions from other strengthening mechanisms assessed to be

small. The transition to the high-temperature, diffusion-controlled regime occurs in these alloys in a rather wide temperature interval of $\sim 0.56\text{--}0.7 T_m$ (Figure 7b). Because the reported multi-phase RCCAs have relatively low T_m , not exceeding 1780°C , the effect from solid solution strengthening is lost at temperatures considerably lower than in many single-phase RCCAs with higher T_m . For example, $0.6 T_m$ represents temperatures below 1000°C for all the studied multi-phase RCCAs. However, precipitation and dispersoid strengthening can be noticeable at temperatures as high as $\sim 0.7 T_m$ for alloys that have $T_s \geq 0.75 T_m$ or which are multi-phase at T_m . The high-temperature strength of these RCCAs is generally lost when the volume fraction of the secondary phases becomes less than $\sim 20\%$. This analysis suggests that multi-phase RCCAs with high T_s and T_m would be better candidates for high-temperature applications (at $T < T_s$) than single-phase RCCAs with the same T_m .

(a)

(b)

Figure 7. Plots of $\log(\sigma_{0.2})$ vs T/T_m for (a) RCCAs reported to be single-phase BCC at 25°C and (b) RCCAs reported to be multi-phase at 25°C. For comparison, curves are shown in figure 7(a) for W [29] and for commercial refractory alloys TZM (a Mo-based alloy) [30] and C-103 (a Nb-based alloy) [31].

4.2 Future Development of High Temperature RCCAs

In the process of improving the properties of RCCAs at high temperatures, two main strategies can be proposed from the above analysis. One strategy is to develop single-phase solid solution RCCAs with high melting temperatures. This approach is based on the observed tendency for the single-phase RCCAs (i) to increase their high-temperature strength with increasing T_m and (ii) to have higher T_m than multi-phase RCCAs. To explore the high T_m single-phase strategy, the variation in T_m values for the vast number of alloying possibilities is estimated using the rule of mixtures (ROM) corrected by an empirical relationship found between T_m^{ROM} and $T_m^{CALPHAD}$, i.e. $T_m = 4(T_m^{ROM} - 807)/3$ (Fig.2). Figure 8 shows T_m as a function of density for *equimolar* mixtures of 455 ternaries, 1372 quaternaries, 3003 quinary and 5005 senary combinations pulled from a panel of 15 elements including Al, Co, Cr, Fe, Hf, Mo, Nb, Ni, Re, Si, Ta, Ti, V, W and Zr. There are two main trends. One is an averaging trend on the barycenter of the scatterplot ($\rho = 9.6 \text{ g/cm}^3$ and $T_m = 1666^\circ\text{C}$) as the number of principal elements increases. The second is the increase of T_m with increasing density. From the 9835 possible equimolar mixtures of principal elements, 1804 can potentially have T_m above 2000°C . These 1804 high melting temperature alloys have densities in the range from 8 to 19 g/cm^3 with a mean value of 12.3

g/cm^3 . Thus, single-phase RCCAs with higher T_m will also have higher alloy density.

Figure 8: Contour lines bounding the melting temperature as a function of density for all possible equimolar combinations of 3, 4, 5 and 6 elements from a palette of 15 elements including Al, Co, Cr, Fe, Hf, Mo, Nb, Ni, Re, Si, Ta, Ti, V, W and Zr. The averaging effect of increasing the number of principal elements shrinks the ranges in densities and melting temperatures toward $\rho = 9.6 \text{ g/cm}^3$ and $T_m = 1666^\circ\text{C}$ (black dot).

Alloy density is an important material constant for alloys used in flight applications and alloys with high specific strength (high strength to density ratio) are preferable for reducing weight and fuel consumption and decreasing environmental pollution. Conventional refractory alloys have high density, generally noticeably higher than other structural metals, contributing to limited use in aerospace structures. On the other hand, recently developed RCCAs have a wide range of densities, varying from 5.6 to 14 g/cm^3 . Higher density single-phase RCCAs generally have higher T_m ; thus, in accord to this work, they should be stronger at high temperatures. It would therefore be interesting to compare the high temperature strengths of RCCAs having different densities and to identify how alloy density affects the specific yield strength of RCCAs at different temperatures.

Figure 9a shows $\sigma_{0.2}$ at 25°C of single-phase and multi-phase RCCAs as a function of alloy density. Only single-phase alloys have densities of 10 g/cm^3 or higher. The peak $\sigma_{0.2}^{25^\circ\text{C}}$ values, exceeding 2000 MPa, are reported for multi-phase RCCAs having densities of $\sim 7\text{-}8 \text{ g/cm}^3$. $\sigma_{0.2}^{25^\circ\text{C}}$ decreases rapidly with decreasing ρ below 7 g/cm^3 or with increasing ρ above 8 g/cm^3 ,

approaching $\sigma_{0.2}^{25^\circ\text{C}} \sim 1000$ MPa for both low-density (~ 6 g/cm³) and high density (~ 14 g/cm³) RCCAs. The specific yield strength, $\sigma_{0.2}^{25^\circ\text{C}}/\rho$, is in the range from ~ 150 to 280 MPa cm³/g for low-density (< 8 g/cm³) RCCAs and it decreases almost linearly to $\sigma_{0.2}^{25^\circ\text{C}}/\rho = 70$ MPa cm³/g with increasing ρ to 14 g/cm³.

At 1000°C , RCCAs with densities less than 10 g/cm³ do not show clear dependence of $\sigma_{0.2}^{1000^\circ\text{C}}$ and $\sigma_{0.2}^{1000^\circ\text{C}}/\rho$ on ρ (Figure 9b), although the specific strength of some multi-phase RCCAs seem to be superior to single-phase RCCAs. RCCAs within this density range have $\sigma_{0.2}^{1000^\circ\text{C}}$ from ~ 80 to 800 MPa and $\sigma_{0.2}^{1000^\circ\text{C}}/\rho$ from 11 to 105 MPa cm³/g. With an increase in ρ from 10 to 14 g/cm³, $\sigma_{0.2}^{1000^\circ\text{C}}$ decreases from 845 MPa to 540 MPa and $\sigma_{0.2}^{1000^\circ\text{C}}/\rho$ decreases from 85 to 40 MPa cm³/g. One can see that high specific strength at 25°C and 1000°C is expected from medium density (~ 6.5 - 9 g/cm³) RCCAs.

The behavior at 1200°C is dramatically different (Figure 9c). Here a clear tendency is seen for $\sigma_{0.2}^{1200^\circ\text{C}}$ to increase with increasing ρ until a peak strength of 735 MPa is achieved at $\rho = 12.3$ g/cm³ for MoNbTaVW. The $\sigma_{0.2}^{1200^\circ\text{C}}/\rho$ values for RCCAs with $\rho \geq 10$ g/cm³ are noticeably higher than for the lower density RCCAs at this temperature. This trend is likely related to the observation that single-phase RCCAs with the highest density (e.g. those containing W) also have the highest T_m values and that $\sigma_{0.2}^{1200^\circ\text{C}}$ increases with T_m faster than the density (Table 1, Table 2, Figure 6b).

Figure 9. (a, b, c) Yield strength ($\sigma_{0.2}$) of single-phase (S) and multi-phase (M) RCCAs at (a) 25°C, (b) 1000°C and (c) 1200°C as a function of alloy density. The dashed performance index lines give constant values of the yield strength to density ratio (specific yield strength). The arrow shows the increase in specific yield strength from the bottom-right to the top-left. The corresponding value of the specific yield strength is shown on the first performance line, and the increment of the specific yield strength between the nearest lines is shown on the second line. Materials with higher performance indices have higher values of the specific yield strength.

The second design strategy for high temperature RCCAs relies on developing multi-phase RCCAs with both high T_s and T_m . This approach is based on the observation that below T_s multi-phase RCCAs generally have higher strength than many single-phase RCCAs, even if their melting temperature is lower and $T_s/T_m > 0.6$. The basic idea is to distribute a high volume fraction of a hard ordered phase, which provides the high temperature strength, in a soft disordered solid solution matrix, which provides the room temperature ductility. Exploring this concept is much more difficult than the previous because it requires prediction of the extent of the solvus in composition and temperature space. There is no simple relationship or correlation that can be used to estimate T_s as a function of composition. Accurate thermodynamic descriptions of the Gibbs free energies are needed that are not being met for high order concentrated mixtures of refractory elements mainly because important intermetallic phases, e.g. ordered bcc structures (B2, L2₁ and D0₃), have no models in refractory alloys. While thermodynamic databases are slowly being improved, innovative approaches should be explored to more rapidly overcome the limitations of current CALPHAD databases. To this respect, the diffusion of artificial intelligence (data-mining and machine learning) in the field of thermodynamic, metallurgy and alloy design represents a great opportunity to anticipate the

stability of phases of interests in high dimensionality composition and temperature space, and point out where to focus experimental effort.

SUMMARY AND CONCLUSIONS

The objective of the present work is to explore origins of the temperature dependent decrease in the yield strength of refractory complex concentrated alloys (RCCAs) at elevated temperatures. Correlations are considered between compressive yield strengths and microstructure (single-phase alloys are compared with multi-phase alloys), alloy melting temperature (taken as the solidus temperature, T_m), solvus temperature (T_s) and volume fraction of secondary phases (V), and an atom misfit parameter (δ_f) in multi-phase RCCAs.

The reported single-phase RCCAs show a wide spread in T_m ranging from 1632°C for $\text{Al}_{0.4}\text{Hf}_{0.6}\text{NbTaTiZr}$ to 2807°C for MoNbTaW . The T_m range of multi-phase RCCAs is much narrower, from 1397°C for $\text{CrMo}_{0.5}\text{NbTa}_{0.5}\text{TiZr}$ to 1780°C for AlCrMoNbTi . Formation of secondary phases during solidification is associated with lower T_m values.

The RT strengths of single-phase and multi-phase RCCAs with volume fractions V of secondary phases less than 40%, are similar and in the range from 880 MPa to 1790 MPa. No clear relationship was found between $\sigma_{0.2}^{25^\circ\text{C}}$ and T_m or δ_f for these alloys. The $\sigma_{0.2}^{25^\circ\text{C}}$ of multi-phase RCCAs tends to increase with increasing V , and alloys with $V \geq 40\%$ are generally stronger than single-phase RCCAs.

Single-phase and multi-phase RCCAs have a similar spread in $\sigma_{0.2}$, from 80 MPa to 855 MPa, at 1000°C. $\sigma_{0.2}^{1000^\circ\text{C}}$ of single-phase RCCAs has a tendency to increase with increasing T_m . $\sigma_{0.2}^{1000^\circ\text{C}}$ of the multi-phase RCCAs shows no dependence on T_m , but it depends on the type of secondary phase and increases with increasing total volume fraction of secondary phases. An increase in T_s of the same type of the secondary phase beyond 1000°C increases $\sigma_{0.2}^{1000^\circ\text{C}}$.

The drop in strength between 25°C and 1000°C is more dramatic for the single-phase RCCAs with lower T_m and for multi-phase RCCAs with low volume fraction ($<20\%$) of secondary phases. Single-phase RCCAs with $T_m > 2000^\circ\text{C}$, or multi-phase RCCAs with $T_s > 1100^\circ\text{C}$ and $V \geq 20\%$ at 1000°C, have better ability to retain high strength at 1000°C.

There is a general trend for $\sigma_{0.2}^{1200^\circ\text{C}}$, $\sigma_{0.2}^{1200^\circ\text{C}}/\rho$ and $\sigma_{0.2}^{1200^\circ\text{C}}/\sigma_{0.2}^{1000^\circ\text{C}}$ of RCCAs that are single-phase structures at 1200°C to increase with increasing T_m . Multi-phase RCCAs have considerably lower $\sigma_{0.2}^{1200^\circ\text{C}}$ and $\sigma_{0.2}^{1200^\circ\text{C}}/\rho$ than many single-phase RCCAs. Multi-phase

RCCAs do not show any dependence of $\sigma_{0.2}^{1200^{\circ}\text{C}}$ on T_m . Their $\sigma_{0.2}^{1200^{\circ}\text{C}} / \sigma_{0.2}^{1000^{\circ}\text{C}}$ ratio decreases with a decrease in the volume fraction of secondary phases.

The analysis of the temperature dependence of the yield strength suggests that solid-solution strengthening in single-phase BCC RCCAs changes from thermally activated dislocation glide below $0.5\text{-}0.6T_m$ to diffusion-controlled dislocation climb at higher temperatures, which causes a rapid drop in the strength above $\sim 0.6T_m$. Thus RCCAs with higher T_m experience this drop at higher temperatures.

The strength of the multi-phase RCCAs is controlled by solid-solution, precipitation and dispersoid strengthening. As the reported multi-phase RCCAs have relatively low T_m , not exceeding 1780°C , the effect from solid solution strengthening is lost at temperatures considerably smaller than for many single-phase RCCAs with higher T_m . However, precipitation and dispersoid strengthening can be noticeable at temperatures as high as $0.7T_m$. The high-temperature strength of these RCCAs is generally lost when the volume fraction of the secondary phases becomes less than $\sim 20\%$. This analysis suggests that multi-phase RCCAs with high T_s and T_m would be better candidates for high-temperature applications (at $T < T_s$) than single-phase RCCAs with the same T_m .

Single-phase RCCAs with higher T_m generally have higher density. Fortunately, their high-temperature strength (at $T = 1200^{\circ}\text{C}$) has the tendency to increase with increasing T_m faster than their density. Therefore, the specific strength of the single-phase RCCAs at 1200°C increases with increasing density or T_m . It is likely that, in order to achieve high specific strength in extreme high-temperature applications, future alloys will have high density, exceeding 10 g/cm^3 . The conducted analysis suggests that the loss of high-temperature strength of single-phase BCC RCCAs is likely related to the activation of diffusion-controlled deformation mechanisms, which occurs at $T \geq 0.6$ of the absolute melting temperature T_m , and that alloys with higher T_m retain their strength to higher temperatures. On the other hand, a rapid decrease in strength of multi-phase RCCAs with increasing temperature above 1000°C is probably due to dissolution of secondary phases, which is controlled by the solvus temperature T_s of the phase responsible for precipitation strengthening. The rapid strength drop of these multi-phase alloys above T_s , where they become single-phase solid solutions, is related to their T_m , similar to single-phase alloys.

ACKNOWLEDGEMENTS

Extensive discussion with Dr. S.I. Rao is greatly appreciated. Work by ONS was supported through the Air Force on-site contract FA8650-15-D-5230 managed by UES, Inc., Dayton, Ohio.

REFERENCES

- [1] Senkov ON, Miracle DB, Chaput KJ, Couzinie J-P. *Journal of Materials Research* 2018;33:3092.
- [2] Gorsse S, Miracle DB, Senkov ON. *Acta Mater.* 2017;135:177.
- [3] Gorsse S, Couzinié J-P, Miracle DB. *C. R. Physique* 2018;19:721.
- [4] Senkov ON, Wilks GB, Miracle DB, Chuang CP, Liaw PK. *Intermetallics* 2010;18:1758.
- [5] Senkov ON, Wilks GB, Scott JM, Miracle DB. *Intermetallics* 2011;19:698.
- [6] Couzinié J-P, Senkov ON, Miracle DB, Dirras G. *Data in Brief* 2018;21:1622.
- [7] Gorsse S, Nguyen MH, Senkov ON, Miracle DB. *Data in Brief* 2018;21:2664.
- [8] Cordero ZC, Knight BE, Schuh CA. *Int. Mater. Rev.* 2016;61:495.
- [9] Yoshida S, Ikeuchi T, Bhattacharjee T, Bai Y, Shibata A, Tsuji N. *Acta Mater.* 2019;171:201.
- [10] Chen S, Tseng KK, Tong Y, Li W, Tsai CW, Yeh JW, Liaw PK. *J. Alloys Cmpds* 2019;795:19.
- [11] Mao H, Chen H-L, Chen Q. *J. Phase Equilibria and Diffusion* 2017;38:353.
- [12] Chen H-L, Mao H, Chen Q. *Mater. Chem. Phys.* 2018;210:279.
- [13] CompuTherm Software, Databases. vol. 2018.
http://www.computherm.com/index.php?route=product/category&path=59_83.
- [14] Zhang F, Kattner U. *Journal of Phase Equilibria and Diffusion* 2015;36:1.
- [15] Senkov ON, Miller JD, Miracle DB, Woodward C. *Calphad: Computer Coupling of Phase Diagrams and Thermochemistry* 2015;50:32.
- [16] Chou KC, Chang YA. *Ber. Bunsenges. Phys. Chem.* 1989;93:735.
- [17] Wertz K, Miller J, Senkov O. *J. Mater. Res.* 2018;33:3204.
- [18] Gorsse S, Tancrét F. *J. Mater. Res.* 2018;33:2899.
- [19] Gorsse S, Senkov ON. *Entropy* 2018;20:899.
- [20] Senkov ON, Woodward C, Miracle DB. *JOM* 2014;66:2030.
- [21] Juan C-C, Tsai M-H, Tsai C-W, Lin C-M, Wang W-R, Yang C-C, Chen S-K, Lin S-J, Yeh J-W. *Intermetallics* 2015;62:76.
- [22] Han ZD, Chen N, Zhao SF, Fan LW, Yang GN, Shao Y, Yao KF. *Intermetallics* 2017;84:153.
- [23] Porter DA, Easterling KE. *Phase Transformations in Metals and Alloys*. Cheltenham, UK: CRC Press, 1992.
- [24] Rao SI, Antillon E, Woodward C, Akdim B, Parthasarathy TA, O.N.Senkov. *Scripta Mater.* 2019;165:103.
- [25] Rao SI, Akdim B, Antillon E, Woodward C, Parthasarathy TA, Senkov ON. *Acta Mater.* 2019;168:222.
- [26] Senkov ON, Scott JM, Senkova SV, Meisenkothen F, Miracle DB, Woodward CF. *J. Mater. Sci.* 2012;47:4062.
- [27] Kuhlmann-Wilsdorf D. *Mater. Sci. Eng. A* 1989;113:1.
- [28] Caillard D, Martin JL. *Thermally Activated Mechanisms in Crystal Plasticity*. Lausanne,

Switzerland: Elsevier, 2003.

[29] Schmidt FF, Ogden HR. The Engineering Properties of Tungsten and Tungsten alloys, DMIC Report 191. Columbus, OH: Battelle Memorial Institute, 1963.

[30] Molybdenum and Molybdenum Alloys. www.molybdenum.com.

[31] ATI Wah Chang Nb Alloy C-103. vol. 2018.

<http://www.matweb.com/search/datasheet.aspx?MatGUID=ed50a3a07706450590669cedc7784150>: MatWeb.com.