

COMPUTATIONAL DESIGN OF ADVANCED AEROTURBINE MATERIALS: NOBURNIUM CYBERALLOY

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Abstract

Under the AFOSR-MEANS initiative, an optimal combination of electronic-level first-principles calculations and efficient experimentation addresses the accelerated design and development of a new class of Nb-based aeroturbine superalloys.

Research Objectives

The objective of this multi-institutional, multidisciplinary effort is to develop optimal strategies for the integration of theory and efficient experimentation for the accelerated design and development of advanced structural materials, using the example of Nb-based superalloys combining oxidation resistance, creep strength and ductility for future aeroturbine blade applications operating at 1300C and above.

Approach

Based on experience in Ni-based superalloys, the primary microstructural concept is a metallic matrix alloy solution, strengthened by a fine dispersion of low-misfit coherent aluminide precipitates incorporating low diffusivity components for coarsening resistance. The precipitates contribute both creep strength and a source of Al for oxidation resistance via Al_2O_3 passivation. Matrix solution requirements include low O solubility and a high diffusivity ratio of Al to O to maintain passivation.

Progress

Our progress has been summarized in an invited overview paper for a recent international symposium on Nb alloys [1]. Guided by phase stability predictions using the FLAPW method, designed model alloys allowed rapid experimental determination of equilibrium tie-tetrahedra in the quaternary Nb-Pd-Hf-Al system bounding the desired Nb-Pd₂HfAl-PdAl three phase field of interest for coherent aluminide dispersion strengthened alloys [2], and alloys were then prepared demonstrating desired finely dispersed microstructures. Extensive FLAPW calculations then

assessed the solution thermodynamics and molar volume behavior of the multicomponent (Pd,Pt)₂(Hf,Zr)Al Heusler phase for improved control of phase stability and interphase misfit, with predicted component intersolubilities validated by selected diffusion couple experiments. BCC solution thermodynamics has been evaluated via a cluster expansion description using the VASP method [2,3]. Using a simple model of O solubility and a preliminary mobility database for prediction of O and Al diffusivities, a team of students in an undergraduate materials design class [4] integrated our models and databases in the theoretical design of oxidation resistant Nb alloys with a combination of Pd₂HfAl and PdAl phases dispersed in a multicomponent BCC Nb-based matrix. Evaluation of their prototype alloys demonstrated predicted microstructures and a substantial level of protective oxide film formation, although predicted Al₂O₃ passivation was not achieved. The student project won first prize in the 2004 TMS-AIME undergraduate design competition.

Based on the preliminary design evaluations, efforts in the past year have emphasized improvement in the mobility models and databases to more precisely control the transport properties governing protective oxide film formation. As a heuristic device for component selection, ADF cluster calculations have assessed the electronic charge density in the octahedral interstitial sites surrounding substitutional atoms in BCC Nb giving the behavior summarized in Figure 1a. It was anticipated that components increasing the charge density will promote lower O solubility while those giving a dramatic decrease in charge density will promote lower O diffusivity through a solution thermodynamic "trapping" effect. Based on this assessment a series of binary alloys with 10^a/o solute were oxidized at 1300C and O diffusion fields were measured by microhardness profiles. The O diffusivities thus measured are summarized in Figure 1b, giving a pattern which is strikingly similar to the relative charge density predictions of Figure 1a. The components Al, Ti, Hf, Y and Pd are shown to strongly reduce O diffusivity with Al showing an order-of-magnitude reduction. The components Al and Hf were then selected for VASP calculations of Nb-X-O ternary solution thermodynamics summarized in Figure 2. The predicted Hf-O chemical attraction is confirmed. For the case of Al which contracts the BCC Nb lattice, an unrelaxed calculation supports a strong Al-O chemical attraction in the nearest octahedral site, but a structurally relaxed calculation shows an overall repulsive interaction attributed to a dominance of elastic interaction in the contracted lattice. The experimentally observed reduction of O diffusivity by Al is thus tentatively attributed to an effect on atomic mobility rather than solution thermodynamics and calculations exploring this are underway. For substitutional diffusion, available measured atomic mobilities in Nb are supplemented by theoretical predictions through a combination of the semi-empirical model of Neumann and Tuijn (*Solid State Phenomena*, Vol. 88, 2002) and preliminary first-principles calculations by Woodward [1]. These models are combined with diffusion couple measurements to define the effect of 10^a/o binary solute additions on the impurity diffusivity of Al at 1300C summarized in Figure 3. The solutes Cr, Pd, Ti, Hf and W are found to enhance Al diffusivity, with the effect of Cr approaching an order of magnitude. Also shown for comparison is the chemical diffusivity of Al at 10^a/o concentration, showing an enhancement by over 2 orders of magnitude.

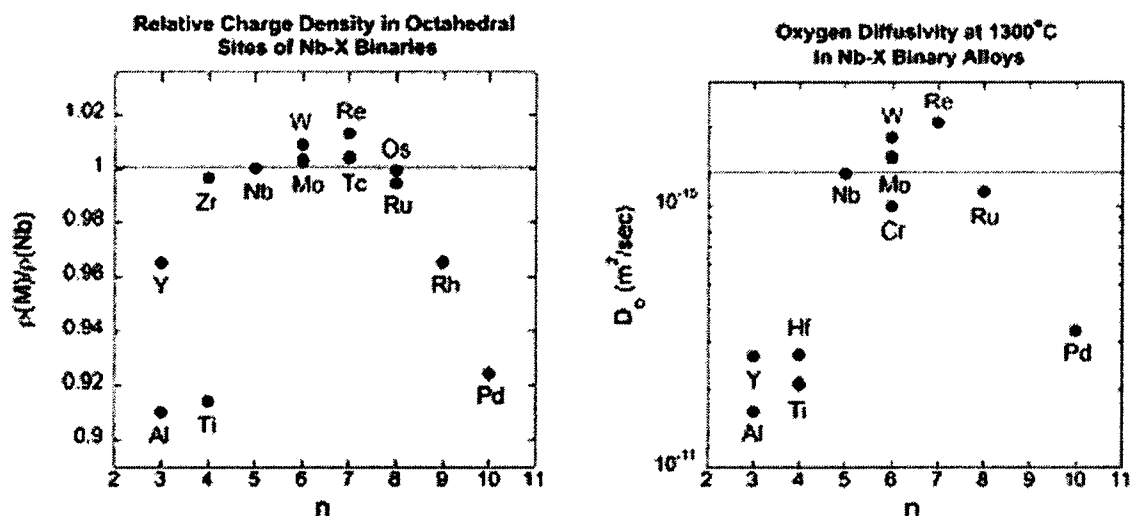


Figure 1 Correlation of calculated relative charge densities in octahedral sites of Nb-X alloys with measured O diffusivities. (a) Calculated charge densities, (b) Measured relative O diffusivities for 10^a/oX at 1300C.

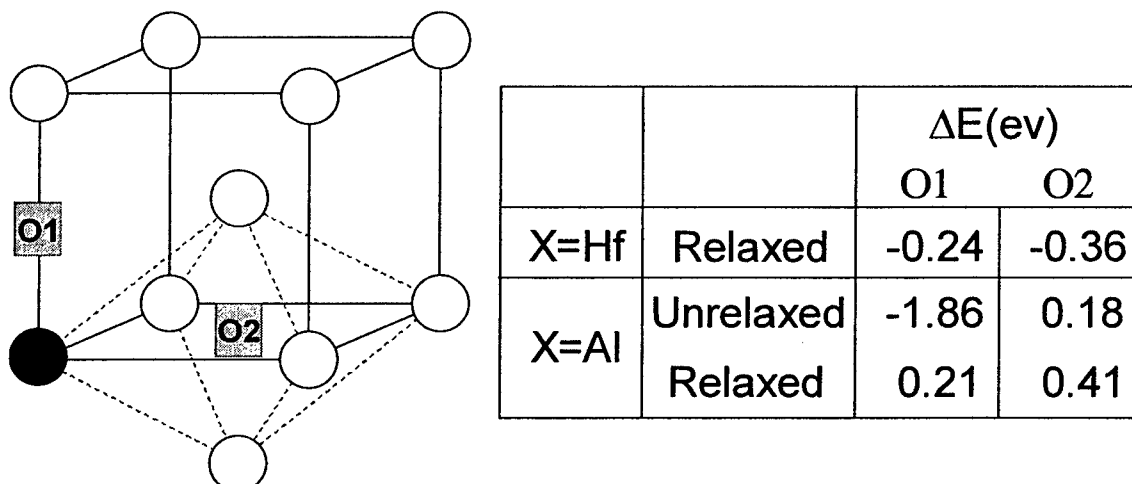


Figure 2 Calculated solute-oxygen binding energies for Hf and Al in BCC Nb.

Guided by the diffusivity behaviors, Al solubilities in BCC Nb at 1300C in Nb-Al-X ternary systems were examined using literature phase diagrams for Cr, Ti and Zr, and our own first-principles calculations for Hf [1]. The latter case shows the highest Al solubility, reaching 21^a/oAl at 34^a/oHf. Preparation of this composition revealed a slightly lower Al solubility of 16^a/o associated with formation of a Nb₂Al dispersion. Although our diffusivity models predict that necessary relative diffusivities of O and Al are satisfied to allow Al₂O₃ passivation, oxidation of the alloy at 1300C produced the complex oxide scale shown in Figure 4 where an unusually compact and adherent (Nb_{0.75}Al_{0.17}Hf_{0.06})₂O₅ external layer forms above a parabolically-growing two-phase rutile-type scale composed of NbAlO₄ and (Hf,Nb,Al)O₂. We

conclude that the desired Al_2O_3 formation is in this case thermodynamically prevented by an unanticipated high stability of the NbAlO_4 oxide. Nonetheless the complex scale shows a parabolic growth constant of $k_p=0.12\text{mg}^2/\text{cm}^4\text{-sec}$ which meets the recession rate over 2000 hrs corresponding to the IHPTET Phase III goals for a 1300C aeroturbine blade alloy. Investigation of the 1300C oxidation behavior of thermodynamically compatible protective aluminide bondcoats for such an alloy show a lower parabolic rate constant by 4.3X for Pd_2HfAl associated with formation of an adherent $\text{Al}_2\text{O}_3+\text{HfO}_2+\text{PdO}_2$ oxide layer, and demonstrate compact Al_2O_3 formation on PdAl .

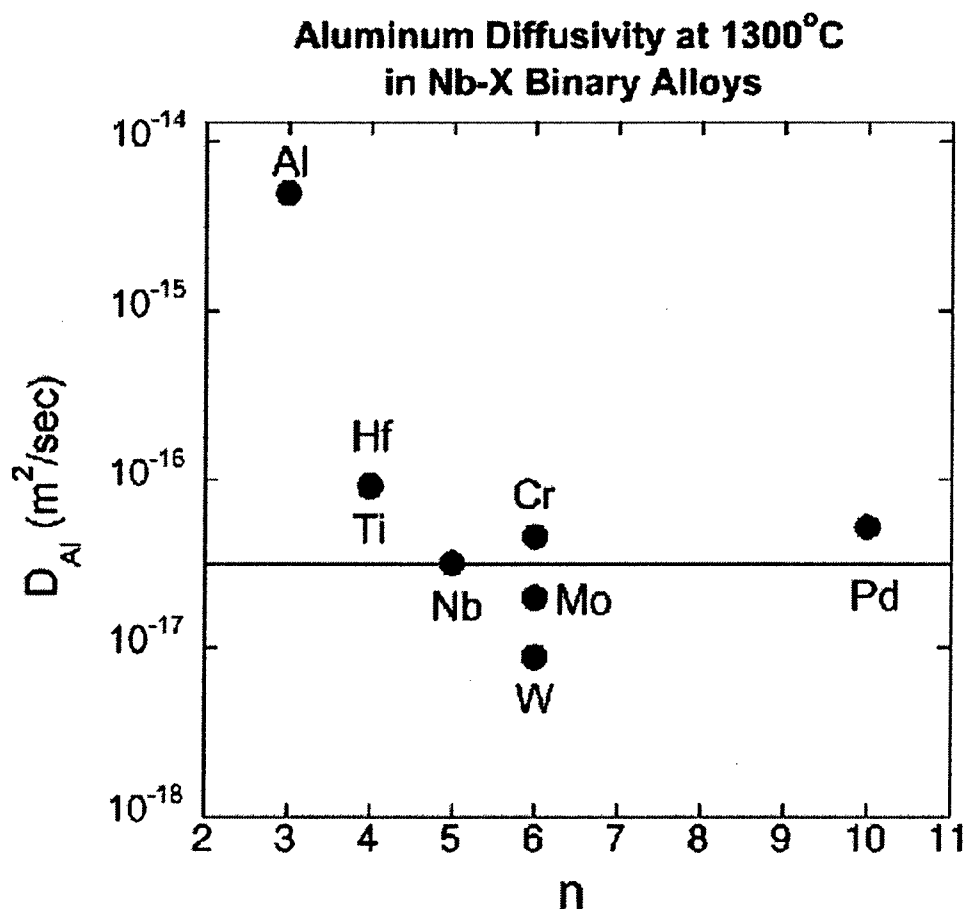


Figure 3 Measured Al impurity diffusivity at 1300C for Nb-X alloys referenced to $10^2/\text{oX}$. Al point represents Al chemical diffusivity at $10^2/\text{oAl}$.

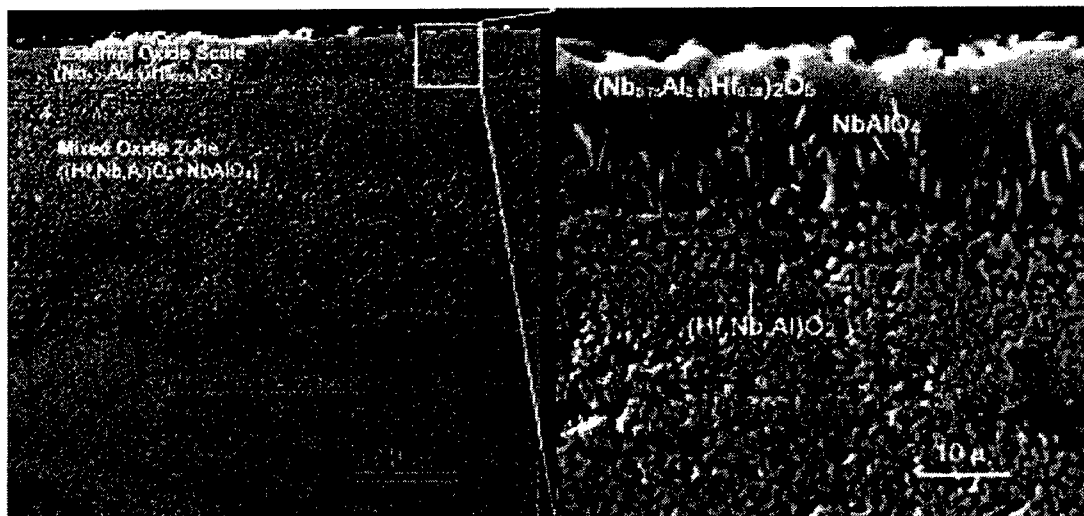


Figure 4 Oxide scale formed on two-phase BCC+Nb₂Al Nb-34a/oHf-21a/oAl at 1300C. Adherent multiphase scale gives an oxidation parabolic rate constant of $k_p=0.12 \text{ mg}^2/\text{cm}^4\text{-sec}$.

To further enhance intrinsic alloy oxidation resistance, a broader survey has been conducted of fast-diffusing components which can promote slow growing oxides of sufficient thermodynamic stability to displace the competing NbAlO₄. Supporting candidate intermetallic phases thermodynamically compatible with Nb have been identified by FLAPW stability calculations, compound melting point measurements and validated phase relations. Oxidation studies of promising alloy prototypes are underway, with preliminary results confirming predicted oxide films with an associated 2 orders of magnitude reduction in k_p .

Acknowledgment/Disclaimer

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Publications

1. G. B. Olson, A. J. Freeman, P. W. Voorhees, G. Ghosh, J. Perepezko, M. Eberhart, and C. Woodward, "Quest for Noburnium: 1300C Cyberalloy," *International Symposium on Niobium for High Temperature Applications*, eds. Y-W. Kim and T. Carneiro, (Warrendale PA, TMS, 2004) 113-122.
2. A. Misra, R. Bishop, G. Ghosh and G.B. Olson, "Phase Relations in Prototype Nb-Pd-Hf-Al Alloys," *Metallurgical and Materials Transactions A*, 34A (2003), 1771-1781.
3. G. Ghosh, A. van de Walle, M. Asta and G. B. Olson, "Phase Stability of the Hf-Nb System: From First-Principles to CALPHAD," *CALPHAD*, 26 (4) (2002), 491-511.
4. E. Altinoglu, J. Bolos and N. Colligan, "Noburnium: Oxidation Resistant Nb Superalloy," Northwestern University, MSc 390 Materials Design project report (May 2003); 2004 TMS-AIME Undergraduate Design Competition winner.

Awards Received

2004 TMS-AIME Undergraduate Design Competition, First Place – N. Altinoglu, J. Bolos and N. Colligan; Project Titled: "Noburnium: Oxidation Resistant Nb Superalloy."

2004 SMD/TMS-AIME Distinguished Scientist/Engineer Award – G. B. Olson.

Transitions

Based on the success of this research, our materials design spinoff company, QuesTek Innovations LLC, has initiated a NASA-sponsored project on design of Nb based alloys for deep space exploration applications.

APPENDIX A

ATOMIC SELECTION FOR ENHANCED OXIDATION RESISTANCE

To overcome the thermodynamic stability of the competing rutile-type NbAlO_4 oxide while satisfying kinetic constraints, thermodynamic and kinetic atomic selection criteria were devised for a broader survey of components and phases for enhanced oxidation resistance of Nb based alloys. The four-dimensional survey is summarized by the Ashby-type selection cross plots in Figure A1. The plot at the left represents the equilibrium thermodynamic parameters of oxide free energy of formation and Pilling-Bedworth (P-B) ratio relative to Nb, while the plot at the right represents 1300C kinetic parameters of oxide scale parabolic rate constant and relative diffusivity in Nb of the corresponding metal solute. Preferred properties in the left plot lie at the upper left, while preferred properties in the right plot lie at the lower right. Using a lower bound estimate of the ΔG_f for NbAlO_4 based on its component Al_2O_3 , and Nb_2O_5 oxides, the thermodynamic plot at the left shows that Y_2O_3 offers desired high stability with the same P-B ratio as the rutile oxides, and mixed Al_2O_3 - Y_2O_3 oxides such as YAG (Yttrium-Aluminum Garnet) offer a good compromise between stability and P-B ratio. Bracketing relative component diffusivities of mixed oxides by their corresponding atomic constituents, the kinetic plot at the right shows that Y_2O_3 also offers the advantage of the high diffusivity of Y in Nb while providing a low k_p equivalent to Cr_2O_3 . The well known low transport properties of YAG offer promise of exceptionally low k_p , but the relative diffusivity will be compromised by the slower diffusion of Al. These considerations identify Y_2O_3 and YAG as preferred oxide scales with the prospect of alloys that could take advantage of the high diffusivity of Y to initially passivate with Y_2O_3 followed by longer time passivation by slower growing YAG.

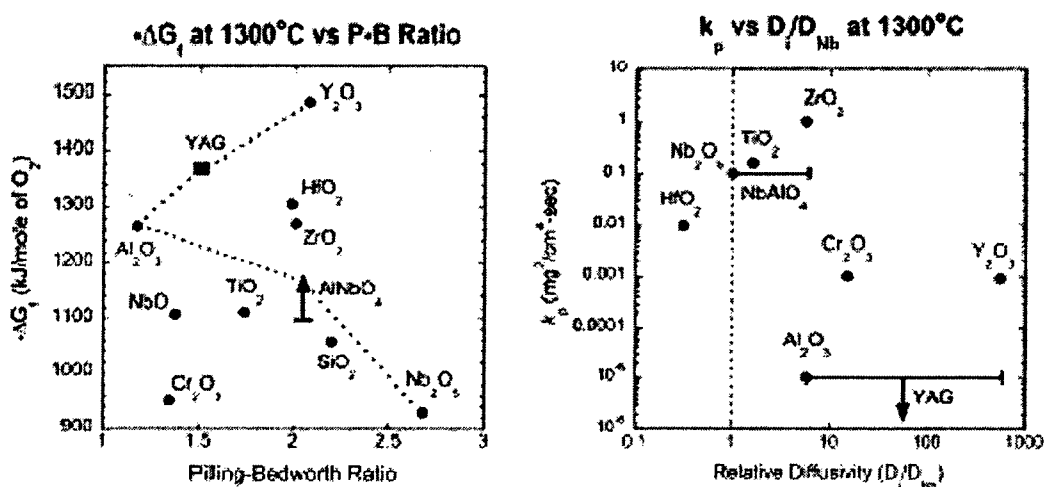


Figure A1 Atomic Selection Parametric Cross Plots.

A challenge to the incorporation of Y in our alloys is the low solubility of Y in Nb and the relatively low melting point of Y compounds. Of the binary Y-Al compounds, Al_2Y offers the highest melting point (1485C) and the nearest stoichiometry to that of YAG. Our experiments show however that it reacts with Nb to form a Nb aluminide and the lower melting point AlY . A literature survey of Y compounds identified those with Pt group metals to be the most stable. Promising Y-Al ternary compounds were then fabricated and their melting points measured by high-temperature DTA. As summarized in Table 1 the equiatomic PtYAl is uniquely distinguished by its melting point above 1500C meeting a primary requirement for our high temperature alloys. Table 2 summarizes our FLAPW calculation of its heat of formation which is greater than -100kJ/mol-atom, exceeding that of our previously studied Heusler compounds. Consistent with this high stability, phase relations measured in model alloys confirm compatibility with Nb. Incorporation into design of a multicomponent alloy optimized for necessary transport properties gives the 1300C microstructure shown at the left in Figure A2, where PtYAl is the predominant dispersed phase. Oxidation at 1300C confirms formation of the predicted stable Y_2O_3 outer scale shown at the right. The scale thickness corresponds to a reduction in oxidation rate constant k_p by two orders of magnitude. Below the Y_2O_3 scale is a mixed layer of $\text{Al}_2\text{O}_3+\text{Y}_2\text{O}_3$ with a dispersion of Pt. Longer time oxidation behavior and the kinetic conditions for conversion of the $\text{Al}_2\text{O}_3+\text{Y}_2\text{O}_3$ layer to the desired stable YAG are under investigation. Our successful predictive design of a Y_2O_3 passivating metallic matrix alloy represents a significant breakthrough in the development of oxidation resistant Nb-based superalloys.

Table 1. Measured melting points of ternary Y-Al compounds

Phase	Crystal Structure	Melting Point
PtYAl	TiNiSi	1580C
PdYAl	TiNiSi	1160C
Pt_2YAl	2 Phase	1120C
Pd_2YAl	Na_3As	890C

Table 2. Computed Compound Heats of Formation-FLAPW

Compound	Structure	Method	ΔH_f , kJ/mol-atom
PtYAl	TiNiSi	GGA LDA	-109.85 -111.78
Pt_2HfAl	L2_1	LDA	-94.44
Pt_2ZrAl	L2_1	LDA	-93.47
Pd_2HfAl	L2_1	LDA	-79.98
Pd_2ZrAl	L2_1	LDA	-79.02

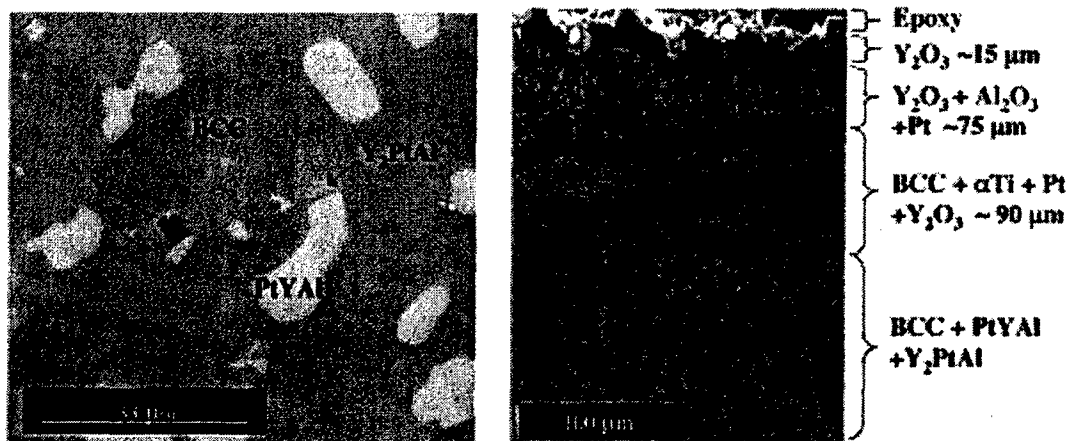


Figure A2 Alloy and Oxide Scale Microstructures in Design Prototype.

QUEST FOR NOBURNIUM: 1300C CYBERALLOY

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Abstract

A multi-institutional, multidisciplinary project addresses optimal integration of computational design and efficient experimentation for the accelerated design and development of high performance materials using the example of Nb-based superalloys combining oxidation resistance, creep strength and ductility for aeroturbine applications operating at 1300°C and above. Integrated within a systems engineering framework, the effort tests the limits of ab-initio quantum mechanical methods to accelerate assessment of thermodynamic and kinetic databases enabling comprehensive predictive design of multicomponent multiphase microstructures as dynamic systems. Based on established principles underlying Ni base superalloys, the central microstructural concept is a dispersion strengthened system in which coherent cubic aluminide phases provide both creep strengthening and a source of Al for Al₂O₃ passivation enabled by a Nb-based BCC alloy matrix with required transport and oxygen solubility behaviors. A combination of FLAPW all-electron and VASP pseudopotential calculations assess thermodynamic and molar volume behaviors of L₂₁ Pd₂HfAl and B2 PdAl based intermetallics as well as BCC Nb-based solutions. Toward an optimal balance of phase stability and interphase misfit, the L₂₁ phase assessment has been extended to the (Pd,Pt)₂(Hf,Zr,Nb)Al system with validation by diffusion couple experiments. To identify slow diffusing species to enhance dispersion coarsening resistance, quantum methods predict an activation energy for Re diffusion in Nb 50% higher than that for Mo in Nb. Employing the DICTRA system, a multicomponent diffusivity database is developed for substitutional alloying elements and interstitial oxygen to enhance the diffusivity ratio of Al to O for promotion of Al₂O₃ passivation. To minimize O solubility, quantum bond topology calculations show trends in interstitial site charge density supporting a dependence of O chemical potential on solution e/a ratio. Integrating the databases and models thusfar developed, a team of students in an undergraduate Materials Design class has performed theoretical designs of 7 component dispersion strengthened alloys with matrix transport and solubility properties supporting feasibility of Al₂O₃ passivation at 1300°C.

Research Objectives

In support of the US Air Force MEANS (Materials Engineering for Affordable New Systems) program,

the objective of this multidisciplinary collaboration is to develop optimal strategies for the integration of theory and efficient experimentation for the accelerated design and development of advanced structural materials, using the example of Nb-based superalloys combining oxidation resistance, creep strength and ductility for future aeroturbine blade applications operating at 1300°C and above.

Approach

Building on demonstrated successes in computational materials design [1], a central theme of the AF-MEANS initiative is the acceleration of the full materials design, development and qualification cycle. This requires not only the extension of materials design tools to process optimization, scaleup, and control of variation, but also the development of new strategies for more efficient integration of theory and experiment. Our systems engineering approach to materials design is represented by Figure 1, summarizing the process-structure-property connections governing the behavior of a multilevel-structured coated Nb-based superalloy system. While the total system integrates compatible thermal barrier and bondcoat subsystems, our initial research has focused on predictive design of the underlying precipitation-strengthened alloy. The specific tools employed in this effort explore the limits of modern quantum engineering methods based on total energy calculations.

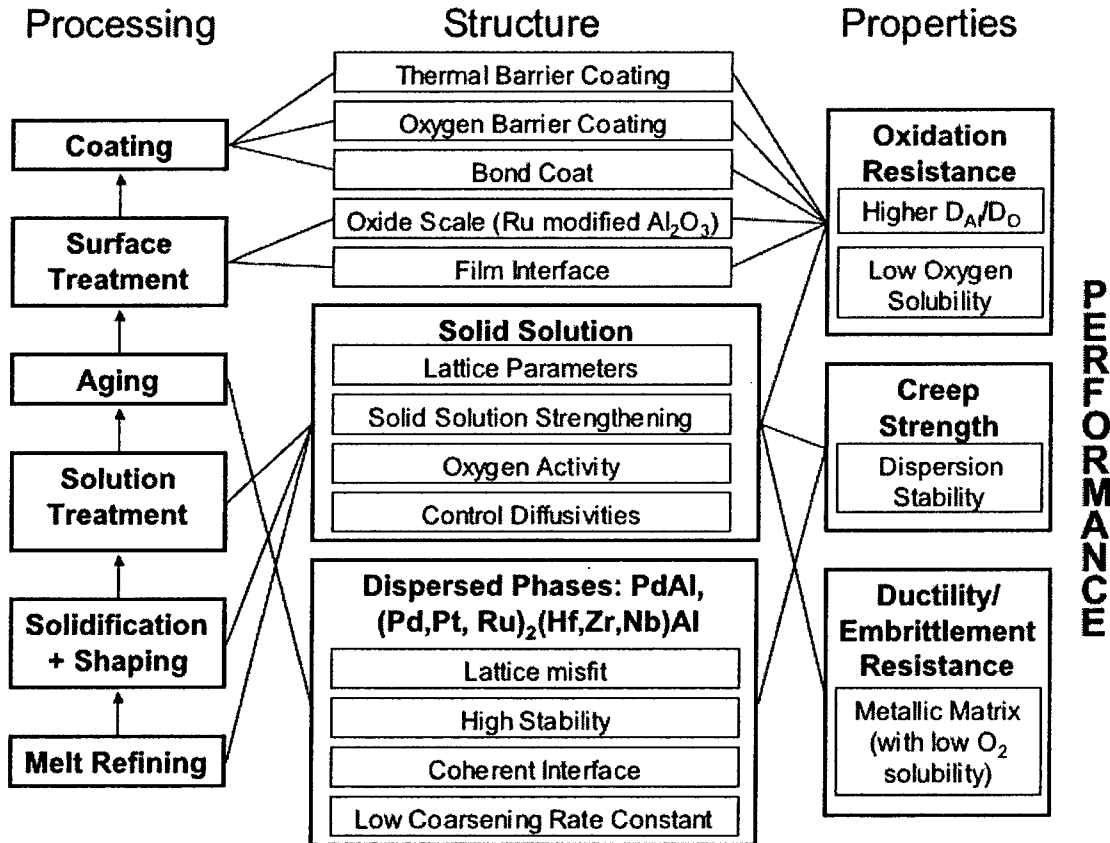


Figure 1. System chart representing coated Nb-based superalloy for aeroturbine blade applications.

Based on experience in Ni-based superalloys, the primary microstructural concept is a metallic matrix alloy solution, strengthened by a fine dispersion of low-misfit coherent aluminide precipitates incorporating low diffusivity components for coarsening resistance. The precipitates contribute both creep strength and a source of Al for oxidation resistance via Al_2O_3 passivation. As predicted by Wagner's analysis [2] the critical alloy Al content, $N_{\text{Al}}^{\text{crit}}$, for formation of a stable passivating Al_2O_3 film is expressed by:

$$N_{\text{Al}}^{\text{crit}} = A \left(\frac{N_{\text{O}}^{(s)} D_{\text{O}}}{D_{\text{Al}}} \right)^{1/2} \quad (1)$$

where A is a constant, $N_{\text{O}}^{(s)}$ is the oxygen solubility in the alloy matrix, and D_{O} and D_{Al} are the diffusivities of oxygen and aluminum, respectively. Matrix solution requirements thus include low O solubility and a high diffusivity ratio of Al to O to maintain passivation.

Results and Discussion

As summarized in Table 1 [3,4], a survey of potential aluminide phases employing available solution thermodynamics and FLMTO quantum-mechanical calculations of compound stabilities identified the Pd_2HfAl L_{21} Heusler phase as the most promising first candidate in terms of stability and lattice parameter, supported by preliminary calculations using the FLAPW method. Quaternary alloys were studied to validate and refine phase relations. Measured tie-tetrahedra in the Nb-Pd-Hf-Al system directly established the composition limits of the desired BCC- Pd_2HfAl two phase field, and a potentially useful BCC- Pd_2HfAl -PdAl three-phase field, as reported in a recent publication [5]. The tie-tetrahedra in the Nb-Pd-Hf-Al system existing at 1200°C and the projection of the tetrahedra on the Pd-Hf-Al basal plane are shown in Figures 2 and 3 respectively.

Table I. Calculated properties for selected Heusler phases [3,4]

Compound	a (calc), nm (at 0 K)	a (expt), nm (at 298 K)	Lattice mismatch (w.r.t pure Nb at 298 K)	$-\Delta E$ (formation) (kJ/mol-atom)
Ni_2TiAl	0.587	0.5872, 0.5865	-11.1% to -11.05%	82.71
Ni_2VAl	0.578	0.633	-12.4% to -4.11%	62.36
Ni_2ZrAl	0.610	0.6123	-7.58% to -7.24%	80.35
Ni_2NbAl	0.600	0.5974	-9.5% to -9.09%	66.83
Ni_2HfAl	0.610	0.6081	-7.88% to -7.58%	91.51
Ni_2TaAl	0.595	0.5949	-9.88% to -9.87%	75.89
Pd_2HfAl	0.633	0.6367	-4.11% to -3.55%	77.69

¹Numbers in parentheses in Figure 3 represent the phase Nb content in at. pct. measured by standard EDS (Energy Dispersive Spectroscopy) SEM analysis using Nb L-alpha Xrays, typically showing less than 1 at. pct. Nb. Recent analysis using Nb K-alpha Xrays however indicates the Nb contents may be as high as 10 at. pct., suggesting high L-alpha absorption, and further microanalysis conducted at a higher accelerating voltage is underway to resolve this apparent discrepancy.

A thorough reassessment of the solution thermodynamics of this system is underway integrating well chosen electronic calculations of compound formation energies employing both FLAPW all-electron and VASP pseudopotential methods. New procedures for efficiently integrating CALPHAD thermodynamic assessments with selective electronic-level total-energy calculations have been developed in collaboration with Prof. Mark Asta of Northwestern using a cluster expansion description of solid solutions. Our recently published assessment of the Nb-Hf system [6] has now been expanded to the Nb-Al and Hf-Al systems to describe the Nb-Hf-Al ternary system (Figure 4). Lattice parameter calculations are integrated into a molar volume database for control of interphase misfit. Efficient thermodynamic assessments combine calculated 0°K formation energies with measured compound melting points and phase boundaries to define entropic contributions.

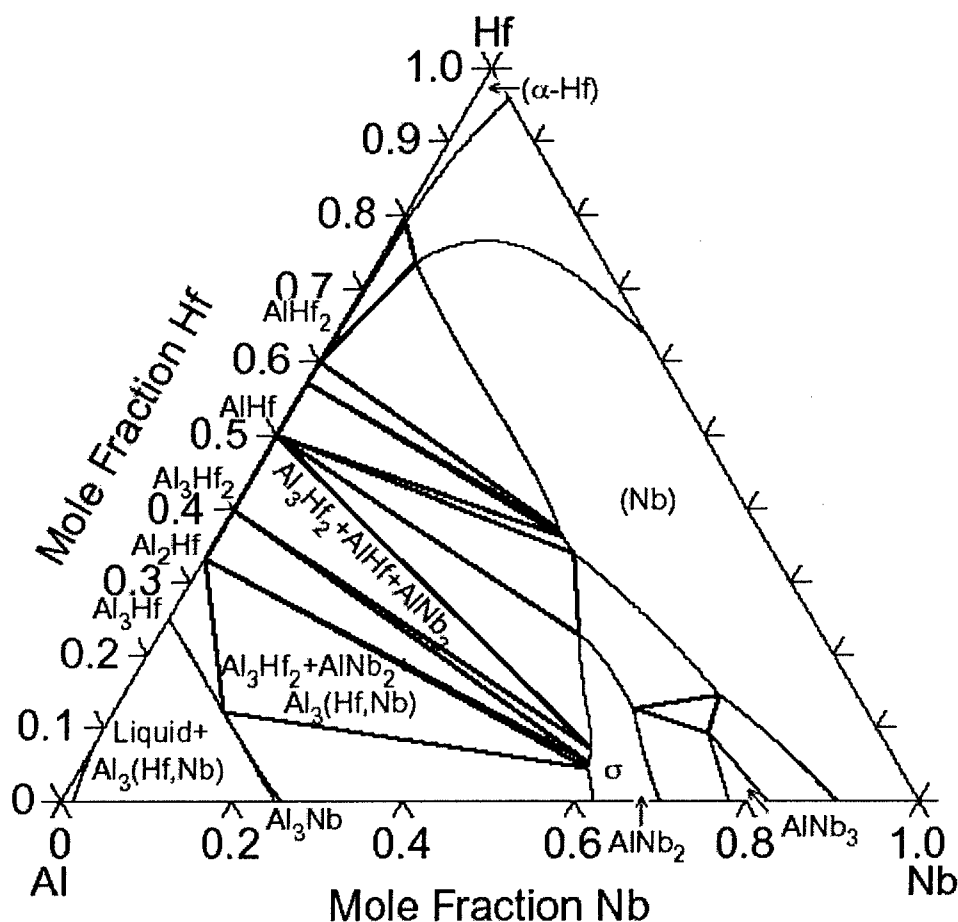


Figure 4. Calculated isothermal section of the Nb-Hf-Al system at 1300°C.

Based on the thermodynamics so far established, model alloys demonstrate desired fine dispersions of L2₁ Pd₂HfAl and B2 PdAl in a Nb matrix as shown by the dark field TEM micrographs of Figure 5 taken using aluminide superlattice reflections [7]. The model alloys show a high microstructural stability at 1000°C compared to Ni-base alloys. Further TEM studies are underway to test for a novel octahedral morphology of coherent precipitates predicted for the unusual "inverse" elastic anisotropy of Nb alloys.

FLAPW calculations for the Heusler phase with partial substitution of Zr in the Hf sublattice predict slight improvement of both thermodynamic stability and lattice parameter for low misfit with Zr addition.

Based on relative stabilities of B2 binary compounds, partial substitution of Pt for Pd was next explored for further improvement. Figure 6 summarizes the predicted enthalpy of formation, enthalpy of mixing, and molar volume for both the Hf-Zr and Pd-Pt intermixing in the $L2_1$ -structured $(\text{Pd,Pt})_2(\text{Hf,Zr})\text{Al}$ Heusler phase. The curves represent best fits of the Redlich-Kister polynomials employed in our solution thermodynamic databases. Although Pt substitution strongly enhances thermodynamic stability and lattice expansion for reduced misfit, the $L2_1$ Heusler phase treated here is no longer the ground state structure in the limit of the $\text{Pt}_2(\text{Hf,Zr})\text{Al}$ compositions. The predicted Hf-Zr intersolubility in $\text{Pd}_2(\text{Hf,Zr})\text{Al}$ has been confirmed by Pd_2HfAl - Pd_2ZrAl diffusion couple experiments at 1100°C (Figure 7). A manuscript reporting these results is in preparation [8].

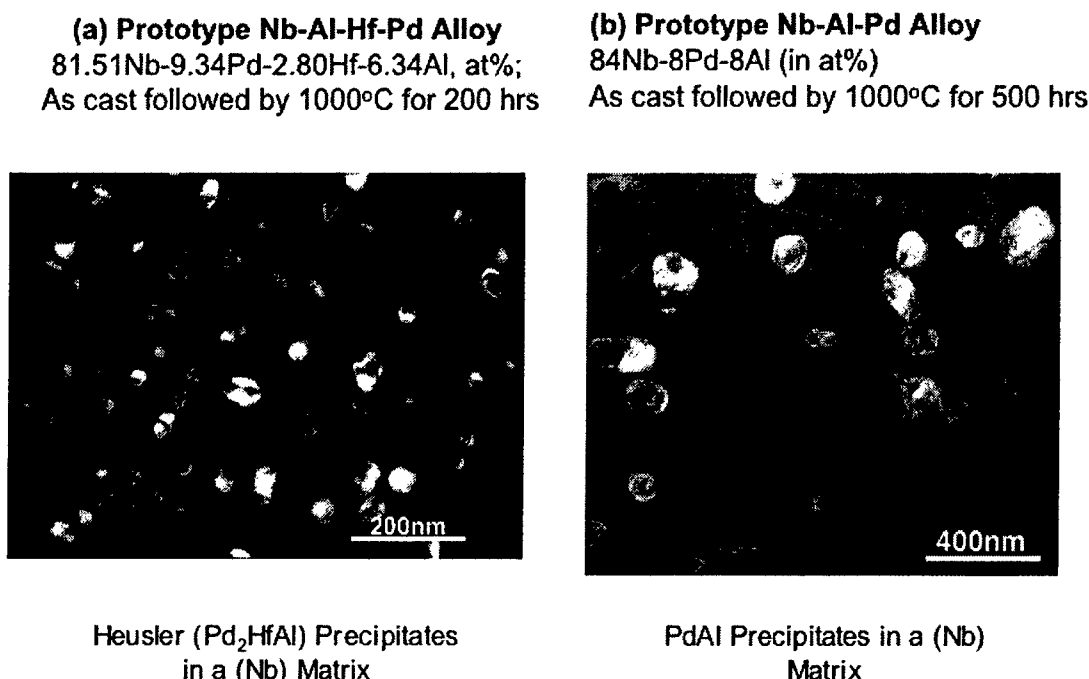


Figure 5. Dark-field TEM micrographs of Pd_2HfAl and PdAl dispersions in Nb alloys [7].

Toward optimization of matrix solution composition, electronic calculations also explore valency effects on interstitial O activity employing bond topological concepts [9]. Computed charge densities in the octahedral sites of BCC Nb and Mo account for an observed [10] dependence of O solubility on alloy electron/atom (e/a) ratio. The predictions for alloying elements in BCC Nb are summarized in Figure 8 showing the trend holds until the e/a of Ru. Short-range order of B2 type is being investigated for the BCC Nb-based solid solution in order to promote 2nd neighbor bonding across the octahedral site as a more efficient means of increasing the site charge density. Incorporating O in both thermodynamic and atomic mobility databases, composition directions are predicted for reduced O solubility and enhanced Al/O diffusivity ratio. The mobility database is also augmented by electronic-level calculations of diffusion energetics with emphasis on prediction of slow-diffusing components for control of multicomponent coarsening resistance [11]. The predicted activation energy for self-diffusion in BCC Ta is in very good agreement with experiment, but is 15% low for the case of Nb. For dilute solution interdiffusivities in Nb alloys, we have employed a five-frequency model for correlation effects incorporating vacancy-solute interactions up to second nearest neighbors. Figure 9 summarizes measured and predicted diffusivities at 1300°C relative to the self-diffusivity of Nb. The predicted relative diffusivity of Mo is in good agreement with experiment. In our search for slow diffusers to enhance microstructural stability, we are very encouraged by the prediction that the activation energy for Re

diffusion is 50% higher than that of Mo, giving a Re diffusivity as low as W, the slowest diffuser yet measured. In support of the prediction, a Nb/Re diffusion couple has shown no measurable intermixing at 1100°C, and higher temperatures are next being investigated.

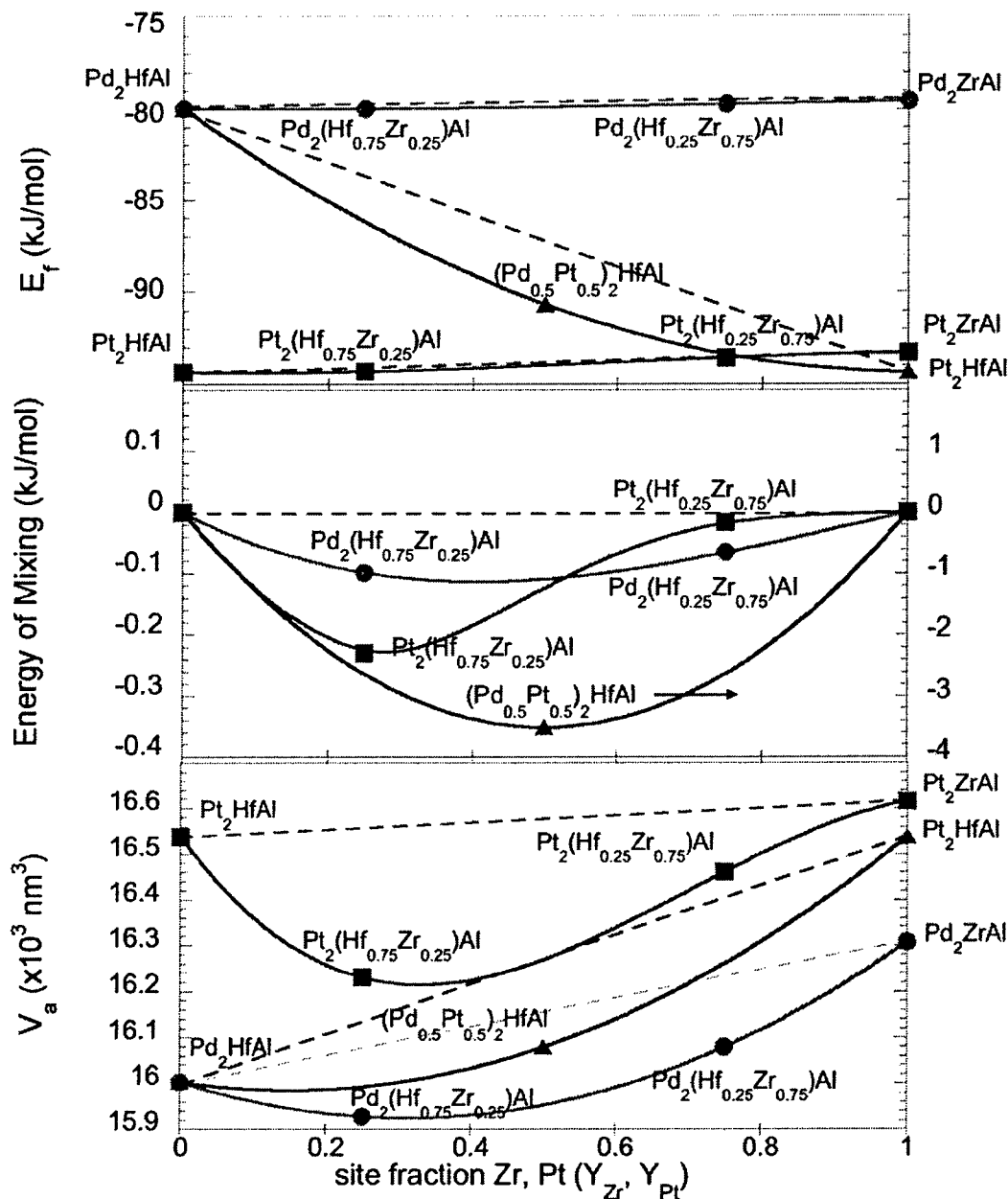


Figure 6. FLAPW calculations of heat of formation, heat of mixing, and molar volume in the $(\text{Pd,Pt})_2(\text{Hf,Zr})\text{Al}$ L_{21} phase.

Preliminary oxidation studies on both model alloys and Heusler intermetallics demonstrate continuous scale formation involving varying combinations of Al_2O_3 and HfO_2 , but do not yet demonstrate desired recession rates. Employing the models and databases thus far developed integrated via the THERMOCALC thermodynamic and DICTRA diffusion kinetic codes, a team of students in an undergraduate Materials Design class at Northwestern has performed the theoretical design of oxidation

resistant Nb alloys with a combination of Pd_2HfAl and PdAl phases dispersed in a multicomponent BCC Nb-based matrix. With thermodynamically compatible additions of Mo, W and Ru in the matrix, the theoretical designs satisfy the Wagner criterion for Al_2O_3 passivation at 1300°C reducing O solubility by 10^{-2} , reducing O diffusivity by 10^{-1} and enhancing Al diffusivity by 10^3 . Their project report [12] was submitted to the TMS-AIME Undergraduate Design Competition, winning 1st Prize for 2004. Prototype alloy validation is underway. Further designs to be evaluated in the next year employ multicomponent oxide scales for more efficient passivation.

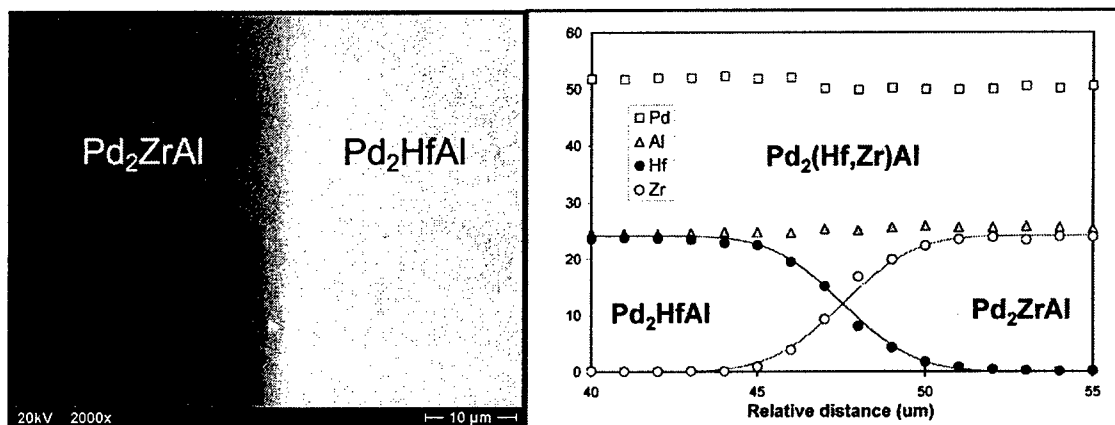


Figure 7. (a) BSE image of cross section of the $\text{Pd}_2\text{ZrAl}/\text{Pd}_2\text{HfAl}$ diffusion couple; (b) Concentration profile obtained from (a) by EPMA [8].

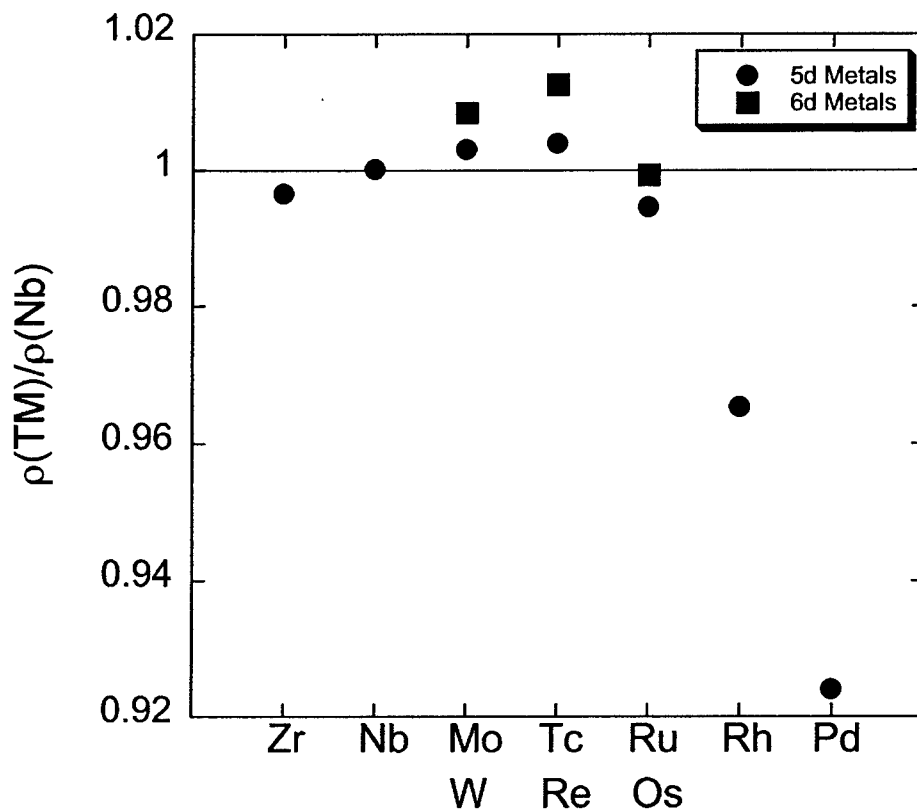


Figure 8. Computed charge density in BCC octahedral sites adjacent to solute atoms in Nb solid solution, scaled to pure Nb case [9].

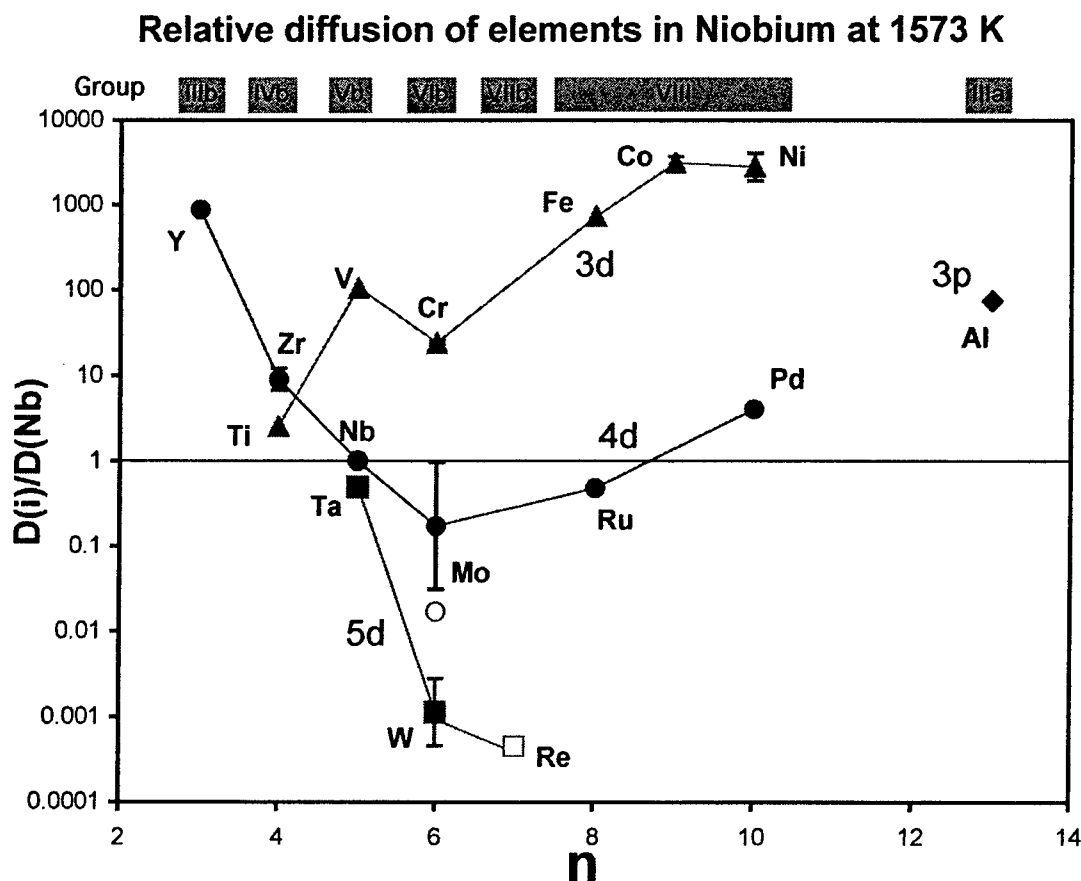


Figure 9. Measured and predicted dilute interdiffusivities in Nb at 1300°C, normalized to Nb self-diffusivity [11].

Conclusions

Efficient integration of quantum mechanical 0°K formation energy calculations with selective measurement of high temperature phase relations allows accelerated development of thermodynamic databases for computational materials design, of particular importance for novel systems such as Nb base alloys where experimental phase diagram data is limited. An important byproduct of the quantum mechanical calculations is the rapid development of molar volume databases supporting control of interphase misfit. Quantitative charge density calculations can also provide guidelines for rapid assessment of interstitial solution thermodynamics governing oxygen solubility behavior. Computational quantum mechanics can also provide useful estimates of substitutional component diffusivities, identifying candidate slow diffusers to enhance microstructural stability at high temperatures.

The integration of these capabilities in the computational design of Nb base alloys exploiting precipitation strengthening by coarsening resistant controlled-misfit coherent multicomponent aluminide phases offers a high potential for combining oxidation resistance and creep strength for application at 1300°C and above.

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APPENDIX C

Final Report for MEANS I

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CURRENT STUDY

Currently, the Ni-based superalloys are used at temperatures close to their operating limits ($< 1100\text{ }^{\circ}\text{C}$) that are greater than 90% of their melting temperatures. Increases in the operating temperature play a key role in advancing in high performance turbine engines because use temperatures limit the efficiency of the turbo engines. In this regard, the Nb-based alloys system has been selected as a high-temperature application because niobium has a high melting temperature ($2468\text{ }^{\circ}\text{C}$) and low density ($8.6 \times 10^3\text{ kg/m}^3$). However, niobium has a poor oxidation resistance. In order to address this issue it is necessary to develop an alloy design that yields a protective oxide scale in the event of an exposure to high temperature environment. Moreover, to minimize high temperature creep, the alloy design includes a strengthening with a fine precipitate of a secondary ordered phase. In the alloy design the $L2_1$ Heusler (X_2YZ) phase has been selected, with aluminum in the Z sites, because the strengthening phase in a given Nb bcc matrix should have a bcc-type crystal structure.

The bcc-type Pd_2XAl ($X = \text{Hf, Zr}$) Heusler phases are attractive since they represent a second phase that exhibits the least lattice parameter discrepancy with the bcc-Nb matrix. Moreover, the formation of stable HfO_2 , ZrO_2 , and Al_2O_3 oxides is expected to play a key role in protecting the multiphase design against oxidation. Nevertheless, the physical property database for the Pd_2XAl Heusler phases is limited. In order to address the high temperature stability of the Nb-based alloys system, it is necessary to examine the high temperature oxidation resistance of the Pd_2XAl Heusler phase and the phase stability between the Pd_2XAl Heusler phase and the bcc-Nb matrix. Therefore, we have facilitated the identification of the key features of the multicomponent phase stability and the oxidation kinetics. The oxidation resistance and stability database for the monolithic Pd_2XAl Heusler phases provide critical input data for the computational design of advanced high temperature materials.

1. $\text{Pd}_2(\text{Hf,Zr})\text{Al}$ Heusler Phases as a second phase

Previously, from a thermodynamic calculation the melting temperature of Pd_2HfAl was estimated at about $3000\text{ }^{\circ}\text{C}$ [bis01]. Based on the binary phase diagram of the Hf-Zr system [asm92], Hf and Zr are completely miscible to form the (αHf , αZr) and (βHf , βZr) solid solution phases, as shown in figure 1. Complete mixing between Hf and Zr leads to predicting a continuous compositional homogeneity region between Pd_2HfAl and Pd_2ZrAl by substitution of Hf to Zr and of Zr to Hf. To confirm the continuity between Pd_2HfAl and Pd_2ZrAl , an $\text{Pd}_2\text{ZrAl}/\text{Pd}_2\text{HfAl}$ diffusion couple was annealed at $1100\text{ }^{\circ}\text{C}$ for 50 hours. A back-scattered electron (BSE) image of the cross section of the annealed $\text{Pd}_2\text{ZrAl}/\text{Pd}_2\text{HfAl}$ diffusion couple is shown in figure 2(a), in which the diffusion reaction zone between Pd_2ZrAl and Pd_2HfAl exhibits a contrast gradient with distance. The continuous contrast change in the diffusion reaction zone indicates the probable continuity of the compositional homogeneity between Pd_2ZrAl and Pd_2HfAl . Concentration profiles (figure 2(b)) obtained from line scanning across

the diffusion reaction zone by an electron probe microanalysis (EPMA) show slight changes in the Pd and Al concentrations between Pd_2ZrAl and Pd_2HfAl , which originally came from difference between nominal composition and fabricated alloy composition of a Heusler phase. The concentrations of Hf and Zr change symmetrically over distance in the diffusion reaction zone. This corroborates that Hf and Zr atoms substitute each other on the X sublattice sites in a Pd_2XAl Heusler phase while Pd and Al atoms remain on their own lattice sites. Consequently, complete mixing of Hf and Zr atoms forms a solution phase $\text{Pd}_2(\text{Hf,Zr})\text{Al}$. The diffusional behavior of Hf and Zr in the $\text{Pd}_2\text{ZrAl}/\text{Pd}_2\text{HfAl}$ diffusion couple can be regarded as a Hf-Zr pseudo-binary interdiffusion in $\text{Pd}_2(\text{Hf,Zr})\text{Al}$. Based on diffusion analysis in a binary system, the interdiffusion coefficient for the Hf-Zr pseudo-binary system in $\text{Pd}_2(\text{Hf,Zr})\text{Al}$ was estimated as $7.9 \times 10^{-18} \text{ m}^2/\text{s}$ at 1100°C .

According to the preliminary pseudo-binary phase diagram calculation of the Nb- Pd_2HfAl system [bis01], the liquidus temperature is $1670 \pm 40^\circ\text{C}$ for Nb-30Pd-15Hf-15Al (all compositions are given as at%) and $1780 \pm 40^\circ\text{C}$ for Nb-20Pd-10Hf-10Al. The pseudo-binary phase diagram also shows that Pd_2HfAl melts congruently above 3000°C . However, the as-cast Pd_2HfAl and Pd_2ZrAl alloys were observed to melt during annealing at 1500°C , which is far below the calculated melting temperature of Pd_2HfAl . A big discrepancy between the melting temperature from the preliminary pseudo-binary phase diagram of the Nb- Pd_2HfAl system and 1500°C at which the as-cast alloys melted, consequently, requires a re-evaluation of the melting temperature of the Pd_2XAl Heusler phase. In order to identify safe temperatures for annealing the $\text{Pd}_2\text{ZrAl}/\text{Pd}_2\text{HfAl}$ diffusion couple, as-cast alloys were heated up with a rate of $40.0^\circ\text{C}/\text{min}$ from 200°C to 1410°C and cooled down with the same rate by a Differential Thermal Analyzer (DTA, Perkin Elmer). The DTA analysis results are shown in figure 3, in which the onset melting temperature of the as-cast Pd_2HfAl and Pd_2ZrAl Heusler phases are estimated at 1440°C and 1350°C . For the as-cast Pd_2HfAl alloy, there exists a small peak at about 1250°C during the cooling process. Identification of the small peak has not yet been completed to determine whether the peak belongs to a solid-state transformation or to solidification of the remaining liquid phase. Continuous homogeneity between Pd_2HfAl and Pd_2ZrAl requires assessing the onset melting temperatures of the $\text{Pd}_2(\text{Hf,Zr})\text{Al}$ phase. Figure 4 shows the assessed onset melting temperatures of the $\text{Pd}_2(\text{Hf}_x\text{Zr}_{1-x})\text{Al}$ where $0 \leq x \leq 1$. The as-cast $\text{Pd}_2(\text{Hf}_x\text{Zr}_{1-x})\text{Al}$ initiates melting at a temperature between 1350 and 1440°C .

2. Oxidation Behavior of the bcc-type Heusler Phases and B2 Phase

A key element of the high temperature Nb alloy design is based upon the use of bcc-related phases in a multiphase microstructure to achieve elevated temperature strengthening and oxidation resistance. To examine oxidation resistance at high temperature, the selected bcc-type Pd_2HfAl (L_{21} Heusler phase), Pd_2ZrAl (L_{21} Heusler phase) and PdAl (B2 CsCl crystal structure) alloys were oxidized at 1100 and 1300°C for 25 hours in air. A cross section SEM-BSE image of the oxidized Pd_2HfAl alloy at 1100°C for 25 hours (figure 5(a)) shows that three distinct layers developed during high temperature oxidation: (i) an outermost dense and hard layer consisting of Al_2O_3 , HfO_2 and PdO_2 , (ii) a mixed layer of oxides (HfO_2 and Al_2O_3) and intermediate phase (PdAl), and (iii) a mixture of PdAl and Pd_3Hf . Composition measurements by an EPMA identified the existence of oxygen in the (i) and (ii) layers, resulting in an oxygen penetration depth about $50 - 60 \mu\text{m}$ out of about $85 - 95 \mu\text{m}$ for the total thickness of three layers. The PdO_2 phase was observed in XRD spectra collected from the oxidized Pd_2HfAl at 1100°C . However, it is reported that PdO_2 decomposes to PdO at 200°C , and PdO decomposes

to Pd at 750 °C [Web]. In this regard, existence of PdO₂ in the oxidized Pd₂HfAl could be interpreted that Pd oxides form during cooling of the sample in air, or that the HfO₂ and Al₂O₃ oxides in the (i) layer limit the decomposition of PdO₂.

The Pd₂HfAl oxidized at 1300 °C for 25 hours developed the (i), (ii) and (iii) layers with a total thickness of about 12-14 μm (figure 5(b)). The oxygen penetration depth at 1300 °C was measured at about 7 - 8 μm. The thickness of three layers formed by the oxidation reaction decreases at higher temperature, indicating an enhancement of oxidation protection with increasing temperature. Figure 6 shows the weight change of Pd₂HfAl as a function of oxidation time that was obtained from an oxidation experiment at 1300 °C by a Thermogravimetric Analyzer (TGA). The TGA analysis indicates the passive oxidation behavior of Pd₂HfAl at 1300 °C because of continuous weight gain. This implies that the inward (from surface to inside) diffusion of oxygen dominates the oxidation kinetics rather than the outward (from inside to surface) diffusion of Pd, Hf and Al. Consequently, oxides form at the interfaces between the (i) and (ii) layers, and the (ii) and (iii) layers during the oxidation reaction.

A microstructural morphology of three distinct layers developed by oxidation reaction can be analyzed to establish an oxidation reaction pathway, (PdO₂+HfO₂+Al₂O₃) / (HfO₂+Al₂O₃+PdAl) / (HfO₂+PdAl) / PdAl / (PdAl+Pd₃Hf) / (PdAl +Pd₃Hf+Pd₂HfAl) / Pd₂HfAl, as shown in figure 5(c). Identification of the three-phase equilibrium among PdAl, Pd₃Hf and Pd₂HfAl in figure 5(c) indicates the stability of the PdAl-Pd₃Hf, Pd₃Hf-Pd₂HfAl and PdAl-Pd₂HfAl two-phase equilibria. Observation of the two-phase equilibria and a three-phase equilibrium coincides with the projected Nb-Pd-Hf-Al tie-tetrahedra bases on the Pd-Hf-Al plane at 1200 °C [mis03]. To determine the phase boundary compositions of PdAl, Pd₃Hf and Pd₂HfAl in their three-phase equilibrium, an EPMA line scanning from the Pd₃Hf-PdAl two-phase equilibrium region to the Pd₂HfAl phase region was performed to obtain the Pd, Hf and Al concentration profiles (figure 5(d)). The phase boundary composition of Pd₂HfAl in the Pd₃Hf-PdAl-Pd₂HfAl three-phase equilibrium that was estimated from the Pd₃Hf/Pd₂HfAl interface in figure 5(d) exhibits a deviation from the stoichiometric composition, indicating an extension of the compositional homogeneity of Pd₂HfAl. According to the concentration profiles in figure 5(d), the composition deviation reveals that the Pd₂HfAl phase is Pd-rich with a composition of Pd_{50+2y}Hf_{25-y}Al_{25-y} (0 ≤ y ≤ 2.5 at%) over a distance of about 30 μm. The formation of the Pd-rich Pd₂HfAl near the Pd₃Hf-PdAl-Pd₂HfAl three-phase equilibrium region (or near the interface between the Pd₂HfAl phase and the PdAl-Pd₃Hf two-phase region) appears to result from Al and Hf transport to three distinct layers that is related to formation of Al₂O₃ and HfO₂ oxides.

A cross section SEM image of the oxidized Pd₂ZrAl alloy at 1100 °C for 25 hours is shown in figure 7, where five distinguishable layers formed during high temperature oxidation: (i) a layer consisting of Al₂O₃, ZrO₂ and PdO₂, (ii) a ZrO₂ oxide layer, (iii) a layer of ZrO₂ and ρ (Pd₇₅Al₂₅), (iv) mixture of Al₂O₃ and ZrO₂, and (v) another layer consisting of ZrO₂ and ρ. Cracks were observed to run through the (ii) layer, resulting in peeling of the oxide layer scale. An EPMA analysis and the SEM image (figure 7) show an oxygen penetration depth of about 700 μm for the 25-hour oxidation. Fast and deep oxygen penetration appears to result in complete consumption of the base Pd₂ZrAl phase because composition measurements by an EPMA identified that the base alloy below the (v) layer exhibited a composition of ρ (Pd₇₅Al₂₅). This indicates that Zr substitution on the X sublattice sites in a Pd₂XAl Heusler phase degrades the oxidation resistance.

The XRD spectra collected from the oxidized PdAl alloys also exhibit the presence of PdO₂. Cross section SEM-BSE images (figure 8) of the oxidized PdAl alloys at 1100 and 1300

°C show that a dense Al_2O_3 oxide layer covers the surface of the PdAl phase and contains Pd. This indicates that the dense Al_2O_3 oxide layer may include PdO_2 that is observed in the XRD analysis. Once again, the existence of PdO_2 in the oxidized PdAl alloys could be interpreted that Pd oxides form during cooling of the sample in air, or that Al_2O_3 limits the decomposition of Pd oxides. The measured Al_2O_3 oxide layer thickness is nearly the same at about 2 μm at 1100 °C and 1300 °C. Consequently, it may be concluded that a thin (1 – 3 μm) oxide layer mixed with Al_2O_3 and PdO_2 protects the PdAl phase at 1100 and 1300 °C, indicating enhancement of the oxidation protection at increasing temperature.

3. Phase Stability and Diffusion Kinetics between Nb and a bcc-type second phase

To examine the high-temperature stability between pure Nb and a Pd_2XAl Heusler phase, diffusion couple experiments were performed. The Nb/ Pd_2ZrAl , Nb/ Pd_2HfAl and Nb/PdAl diffusion couples were annealed at 1100 °C under a Ti-gettered Ar atmosphere. Figure 9(a) shows the SEM-BSE image of the cross section of the Nb/ Pd_2HfAl diffusion couple annealed at 1100 °C for 50 hours. The annealing process generated several intermediate phases between Nb and Pd_2HfAl . Composition measurements by an EPMA determined that the intermediate phases are Hf(ss), PdAl, and Pd_3Hf . Figure 9(a) shows that the Hf(ss) phase includes a fine-scale second phase that is not identified because of its size. During isothermal annealing, the Nb/ Pd_2HfAl diffusion couple developed a diffusion reaction path Nb / Hf(ss) / Hf(ss)+PdAl / PdAl / PdAl+ Pd_3Hf / PdAl+ Pd_3Hf + Pd_2HfAl / Pd_2HfAl at 1100 °C, as shown in figure 9(b). The reaction product layers exhibit a total thickness of about 50 μm .

Figure 10(a) shows a SEM-BSE image of the cross section of the Nb/ Pd_2ZrAl diffusion couple annealed at 1100 °C for 50 hours. The annealing process generated several intermediate phases between Nb and Pd_2ZrAl . The EPMA measurements determined that the intermediate phases are Zr(ss) and PdAl, where the Zr(ss) phase may have some porosity. During isothermal annealing, the Nb/ Pd_2ZrAl diffusion couple developed a diffusion reaction path Nb / Zr(ss) / Zr(ss)+PdAl / Zr(ss)+PdAl+ Pd_2ZrAl / Pd_2ZrAl at 1100 °C, as shown in figure 10(b). The reaction product layers exhibit a total thickness of about 60 μm . In fact, the Hf-Zr binary phase diagram shows formation of the (αHf , αZr) and (βHf , βZr) complete solid solution phases, indicating the similarity of Hf and Zr, and their mutual complete solubility. Consequently, the diffusion kinetics in the Nb/ Pd_2HfAl and Nb/ Pd_2ZrAl couples was expected to be similar. However, during isothermal annealing, the Nb/ Pd_2HfAl diffusion couple developed Hf(ss), PdAl and Pd_3Hf while the Nb/ Pd_2ZrAl diffusion couple developed Zr(ss) and PdAl. This established two distinct diffusion pathways of the Nb/ Pd_2HfAl (figure 9(b)) and Nb/ Pd_2ZrAl (figure 10(b)) couples.

Figure 11(a) shows the SEM-BSE image of a cross section of the Nb/PdAl diffusion couple annealed at 1100 °C for 150 hours. Sluggish Nb, Pd and Al atom movements resulted in a negligible concentration change (figure 11(b)), and no product phase was observed between Nb and PdAl. This result suggests that a Nb-PdAl two-phase equilibrium may exist so that the phase combination of Nb and PdAl at 1100 °C would not be degraded by the formation of byproduct phases. In 1987, Kiseleva calculated the stability of Heusler phases using the relation of the atomic dimension of components and electron concentration as well as thermodynamic and electron properties of components. He predicted stable formation of the Pd_2NbAl Heusler phase. If Pd_2NbAl is in equilibrium with both (Nb) and PdAl, a ternary equilibrium among (Nb), PdAl and Pd_2NbAl would occur (figure 12). This allows for a multiphase design of $\text{BCC}(\text{Nb}) + \text{B2}(\text{PdAl}) + \text{L2}_1(\text{Pd}_2\text{NbAl})$, where L2_1 and B2 belong to the bcc-based crystal structure. Figure

13 shows a SEM-BSE image of as-cast Pd-25Nb-25Al (at% in unit) alloy, in which an EPMA analysis identified the co-existence of (Nb), PdAl and Pd₂NbAl.

To develop an effective database for the diffusion behavior in Nb binary alloys, diffusion experiments have been conducted on interdiffusion in the Nb/Re and Nb/Ru couples. For the diffusion treatment at 1100 °C, the Nb/Re diffusion couple was annealed for 200 hours under a Ti-gettered Ar atmosphere. For the heat treatment at 1600 °C that is about 300 °C higher than the intended use temperature of the Niobium alloys, the Nb/Re diffusion couple was annealed for 100 hours under a high vacuum atmosphere of 7×10^{-7} Torr. The cross section of the 1100 °C diffusion couple is shown in figure 14(a), in which contrast of each end member pure element is discrete at the interface. The Nb and Re concentrations were measured across the interface by an EPMA with a 20kV acceleration voltage and 20 nA current. Based on the Nb and Re concentration profiles, Nb and Re atom movements are sluggish and develop a narrow diffusion zone with a width of about 3 μm along the interface. The profiles elucidate that the diffusion process does not form the χ intermediate phase that exhibits the homogeneity region ranging from 63.5 – 87 at%Re. In other words, Nb atoms diffuse in the (Re) solid solution and Re atoms in the (Nb) solid solution. An error function regression fit on experimentally measured concentration profile data, was used to estimate the interdiffusion coefficient of (Nb) at $\tilde{D} = 2.4 \times 10^{-19} \text{ m}^2/\text{s}$. Because the displacements of the Kirkendall voids (black dots on the interface in figure 14(a)) can be negligible, the estimated interdiffusion coefficient may approximate the intrinsic diffusion coefficient.

The cross section of the Nb/Re diffusion couple annealed at 1600 °C is shown in figure 14(b), in which the χ intermediate phase forms between Nb and Re. An EPMA analysis measured the Nb and Re concentration profiles as a function of distance. According to the profiles, Nb and Re atom movements develop a diffusion reaction zone of about 40 – 45 μm . The (Nb) solution phase exhibits the Re solubility of about 44 at%Re. The compositional homogeneity range of the χ phase extends from 64.3 at%Re to 86.1 at%Re. This is consistent with the published phase diagram of the Nb-Re system, where the Re solubility of (Nb) is 44.3 at% Re and the homogeneity region of χ ranges from 63 – 87 at%Re. By fitting an error function, based on an assumption of constant diffusivity, the estimated interdiffusion coefficients of (Nb) and χ are $2.6 \times 10^{-16} \text{ m}^2/\text{s}$ and $6.0 \times 10^{-18} \text{ m}^2/\text{s}$, respectively. A plot of diffusion coefficients versus the reciprocal temperature is shown in figure 15, in which the interdiffusion coefficients of (Nb) are compared with the Nb self-diffusivities. At 1600 °C the Nb self-diffusivity is faster than the (Nb) interdiffusion coefficient while at 1100 °C the Nb self-diffusivity is slower than the (Nb) interdiffusion coefficient. This means that at 1100 °C the chemical potential gradient for directive atom movements is a dominant driving force rather than the thermal energy for random jumps is.

The Nb/Ru diffusion couple that was annealed at 1200 °C for 200 hours developed two distinct product layers (figure 16(a)); two layers are labeled as layer 1 and layer 2. The Nb and Ru concentration profiles (figure 16(b)) obtained by an EPMA line scanning on the Nb/Ru diffusion couple determined the compositional homogeneity ranges of two layers that are not comparable to a binary Nb-Ru phase diagram (figure 16(c)) in reference **asm92**. This requires a re-assessment of the detailed phase stability in the Nb-Ru binary system. Based on the concentration profiles, as shown in figure 16(b), the interdiffusion coefficient of (Nb) was estimated as $5.0 \times 10^{-18} \text{ m}^2/\text{s}$, which can be compared to the Nb self-diffusivity of $4.3 \times 10^{-19} \text{ m}^2/\text{s}$ at 1200 °C.

SUMMARY OF ACCOMPLISHMENTS

As a selected second phase in the Nb-based alloy design for improving the oxidation resistance and for minimizing high temperature creep, the Pd_2XAl ($\text{X} = \text{Hf}, \text{Zr}$) Heusler phases have been examined. Complete miscibility between Hf and Zr yields a solution phase $\text{Pd}_2(\text{Hf,Zr})\text{Al}$, where Hf and Zr atoms substitute each other on the X sublattice sites in a Pd_2XAl Heusler phase while Pd and Al atoms remain on their own sublattice sites. The DTA analysis reveals that the as cast $\text{Pd}_2(\text{Hf}_x\text{Zr}_{1-x})\text{Al}$ where $0 \leq x \leq 1$ initiates melting at a temperature between 1350 – 1440 °C, which is far below the thermodynamic calculation that Pd_2HfAl melts congruently above 3000 °C.

To examine oxidation resistance, the bcc-type L2_1 Pd_2XAl Heusler phases and B2 PdAl phase were oxidized at 1100 and 1300 °C in air. Three distinct layers developed during oxidation of Pd_2HfAl . The Pd_2HfAl phase below three distinct layers exhibits a Pd-rich composition of $\text{Pd}_{50+2y}\text{Hf}_{25-y}\text{Al}_{25-y}$ ($0 \leq y \leq 2.5$ at%) over a distance of about 30 μm . The TGA analysis indicates the passive oxidation behavior of Pd_2HfAl at 1300 °C because of weight gain. From an analysis of the kinetics it is concluded that the inward diffusion of oxygen dominates the oxidation kinetics. During oxidation of Pd_2ZrAl fast and deep oxygen penetration appeared to result in complete consumption of the base Pd_2ZrAl , indicating that Zr substitution on the X sublattice sites in a Pd_2XAl Heusler phase degrades the oxidation resistance. In the oxidized PdAl alloys it was observed that a dense Al_2O_3 oxide layer covers the surface of PdAl and contains Pd. Measurements of Al_2O_3 oxide layer thickness at 1100 and 1300 °C indicate enhancement of the oxidation protection at increasing temperature.

Diffusion couple experiments have examined the high-temperature phase stability and diffusion kinetics between pure Nb and a bcc-type second phase in the Nb-based alloy design. The Nb/ Pd_2HfAl diffusion couple that was annealed at 1100 °C for 50 hours developed the intermediate phases Hf(ss), PdAl and Pd_3Hf with a total reaction product layers thickness of about 50 μm . The Nb/ Pd_2ZrAl diffusion couple that was annealed at 1100 °C for 50 hours developed the intermediate phases Zr(ss) and PdAl with a total reaction product layers thickness of about 60 μm . Annealing the Nb/PdAl diffusion couple at 1100 °C for 150 hours did not generate any reaction product, indicating a probable Nb-PdAl two-phase equilibrium. This predicts a multiphase design of $\text{BCC}(\text{Nb}) + \text{B2}(\text{PdAl}) + \text{L2}_1(\text{Pd}_2\text{NbAl})$, provided that the Pd_2NbAl Heusler phase exists as a stable phase and is in equilibrium with both (Nb) and PdAl. From the Nb/Re diffusion couple experiments the (Nb) interdiffusion coefficient was estimated as $2.4 \times 10^{-19} \text{ m}^2/\text{sec}$ at 1100 °C and $2.6 \times 10^{-16} \text{ m}^2/\text{sec}$ at 1600 °C. At 1600 °C the Nb self-diffusivity is faster than the (Nb) interdiffusion coefficient while at 1100 °C the Nb self-diffusivity is slower than the (Nb) interdiffusion coefficient. The Nb/Ru diffusion couple that was annealed at 1200 °C for 200 hours developed two distinct product layers. An EPMA analysis shows that the measured compositional homogeneity ranges of two layers are not comparable to the published Nb-Ru binary phase diagram.

FUTURE DIRECTIONS

From the initial observations it is evident that the PdAl phase plays a major role in controlling the oxidation response. The PdAl intermediate phase was observed in the multiple-layer structures that were formed on the Pd_2HfAl and Pd_2ZrAl phases by oxidation. Since PdAl forms an Al_2O_3 oxide layer by oxidation, a Pd-rich PdAl is expected to develop below the Al_2O_3 oxide layer. In fact, the Al_2O_3 oxide layer formed by oxidation reaction at 1300 °C is thinner

than the Al_2O_3 oxide layer formed at 1100 °C, indicating the enhancement of oxidation protection with increasing temperature. Moreover, the crystal structures of B2 PdAl and a L2₁ Heusler phase (Pd_2HfAl , Pd_2ZrAl , etc.) indicate that small additions of Hf or Zr into PdAl may substitute for Al. In this regard, the oxidation behavior could change with addition of Hf or Zr into PdAl and indicate another role for the Hf/Zr ratio in controlling the microstructure. Again, a similar behavior can be expected for PtAl and the substitution of Y for Hf or Zr.

In order to assess oxidation behavior, the PdAl design, in which Pd and Al compositions change continuously with distance, will be subject to heat treatment in air. Al-rich PdAl is expected to be less oxidation resistant behavior than Pd-rich PdAl, provided that PdAl is assumed to exhibit the similar diffusional behavior of NiAl. In the Al-rich PdAl, diffusion of Al atom movements may slow down as the composition below the Al_2O_3 oxide layer reaches stoichiometry. This may increase the oxygen penetration depth and the thickness of the oxide layer. In other words, the oxide layer structure may be expected to change with composition of PdAl.

A number of diffusion couple experiments involving Nb/PdAl, Nb/Re and Nb/Ru pairings should be performed at various temperatures to assess the diffusivities as well as phase stabilities that determine the equilibrium microstructure. Since the standard diffusion couple experiment gives information about phase stability such as a tie line and diffusion kinetic properties only on the diffusion pathway, multiple standard diffusion couple explorations between two phases are necessary. For rapid assessments of phase stability and diffusion kinetics, diffusion couples and diffusion multiples can be employed where the diffusion multiple is an assembly of three or more different alloys. Continued experiments on diffusion couples made by assembling two pure metals such as Nb-Re, Nb-Ru and other pairings can demonstrate the diffusion behavior and contribute essential information for the stable phase boundary database. A diffusion multiple can be constructed by adding another member such as Nb to the $\text{Pd}_2\text{HfAl}/\text{Pd}_2\text{ZrAl}$ couple either before or after Pd_2HfAl and Pd_2ZrAl undergo diffusion reaction. With the addition of the third member the diffusion multiple can be used to probe within multicomponent phase space and identify the multiple tie lines that are necessary for a phase stability analysis. Moreover, based upon the diffusion multiple exploration, specific intermediate phase diffusion couples can be designed to allow for an accurate evaluation of the component diffusivities.

A key objective will be to identify means to control the diffusion pathway. For instance, due to the similarity of Hf and Zr and their mutual solubility, the diffusion kinetics in the Nb/ Pd_2HfAl and Nb/ Pd_2ZrAl couples is expected to be similar. However, following an anneal at 1100 °C, Hf(ss), Pd_3Hf and PdAl developed in the Nb/ Pd_2HfAl diffusion couple while only Zr(ss) and PdAl developed in the Nb/ Pd_2ZrAl diffusion couple. This corroborates that the diffusion behavior between Nb and the $\text{Pd}_2(\text{Hf,Zr})\text{Al}$ phase changes dramatically at certain composition ratio of Hf to Zr. Moreover, the change in the diffusion kinetics between Nb and $\text{Pd}_2(\text{Hf,Zr})\text{Al}$ indicates that the microstructure of the Nb-Pd-(Hf,Zr)-Al system can be controlled by the Hf/Zr composition ratio. Consequently, the Nb/ $\text{Pd}_2(\text{Hf,Zr})\text{Al}$ diffusion multiple experiments with a variation of the Hf/Zr composition ratio are of special interest.

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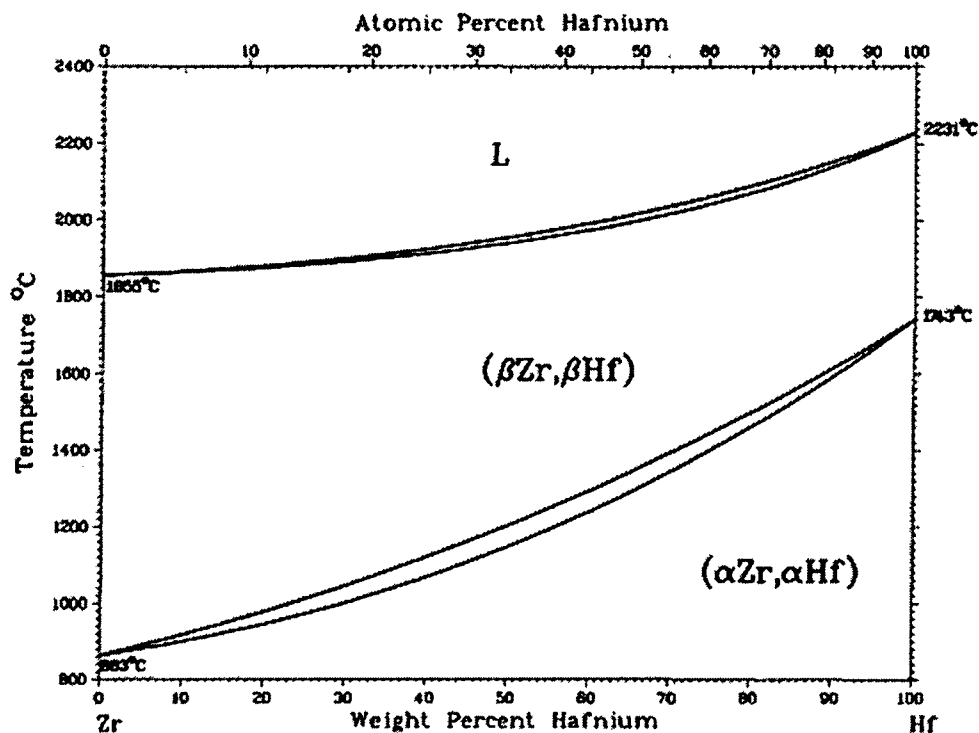


Figure 1. Zr-Hf binary phase diagram [asm]

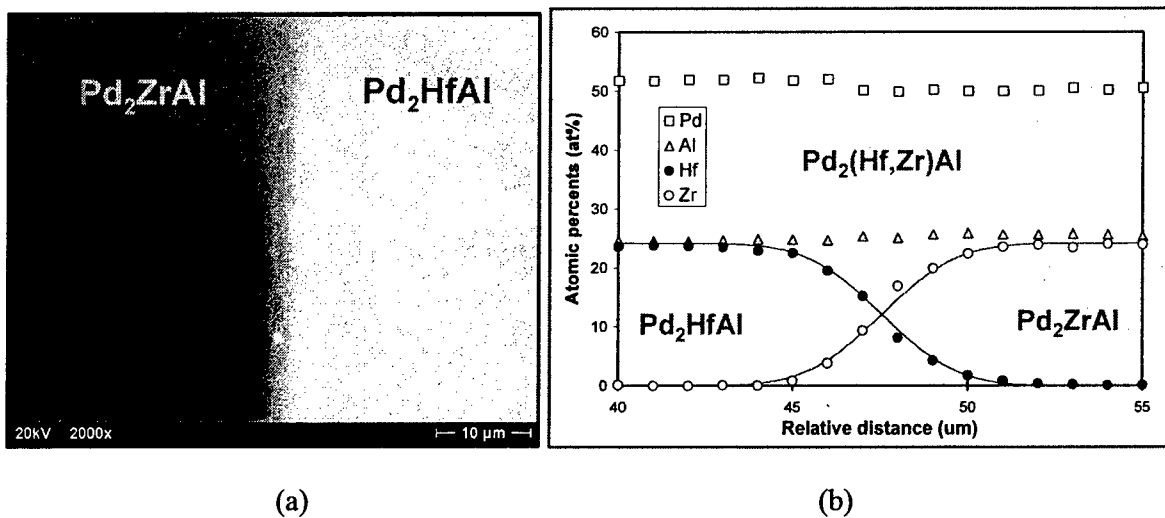


Figure 2. (a) SEM BSE image of the cross-section of the Pd₂HfAl/Pd₂ZrAl diffusion couple annealed at 1100 °C for 50 hours and (b) concentration profiles of Pd, Hf, Zr and Al obtained from the diffusion couple shown in (a).

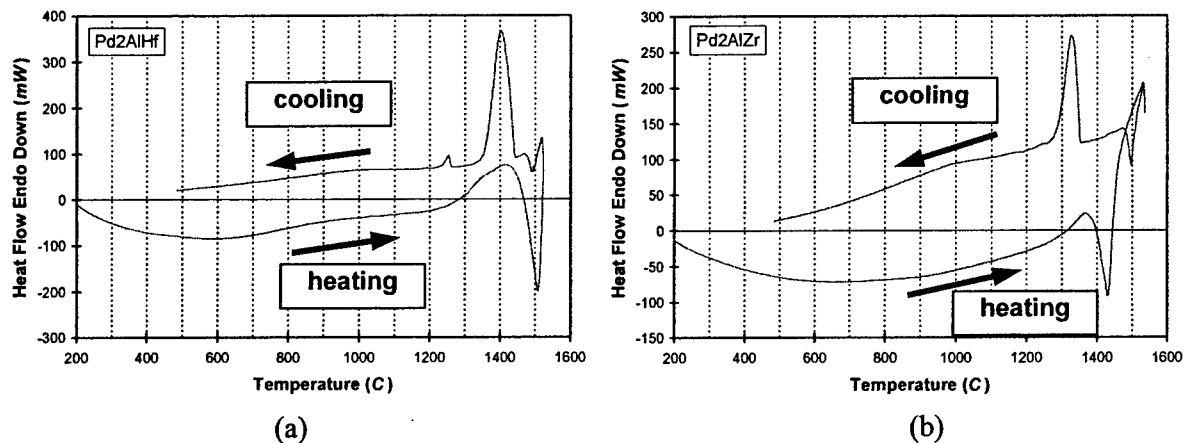


Figure 3. DTA curves of (a) as-cast Pd₂HfAl and (b) as-cast Pd₂ZrAl with a rate of 40.0 °C/min

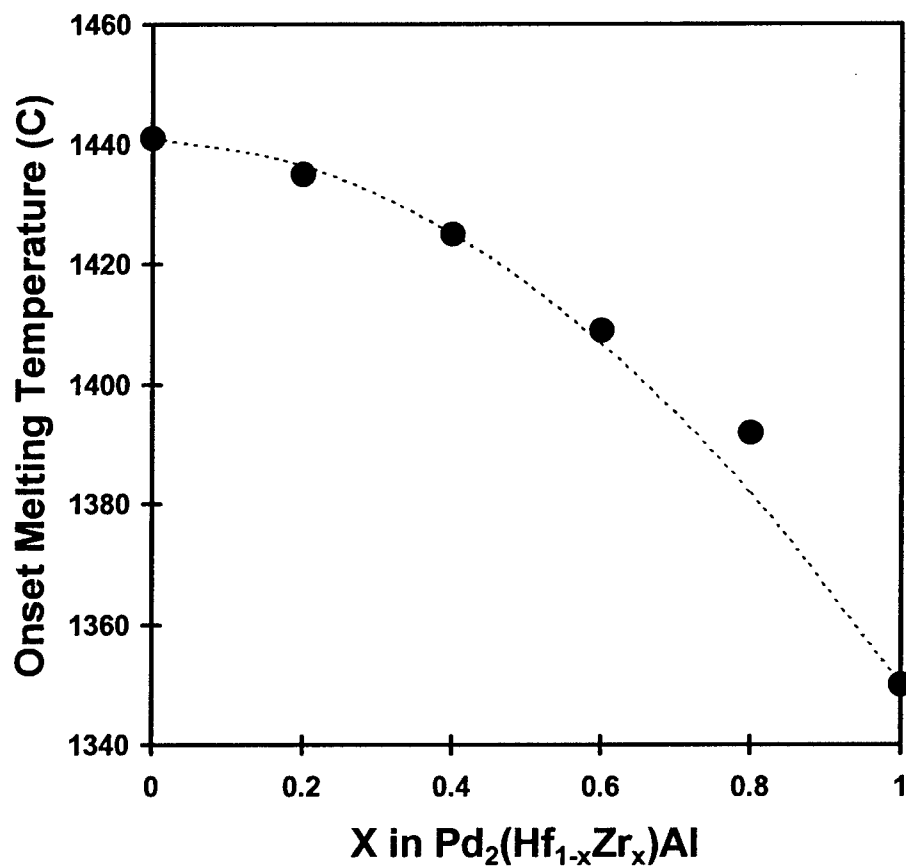


Figure 4. Onset melting temperature of Pd₂(Hf_{1-x}Zr_x)Al, where 0 ≤ x ≤ 1.

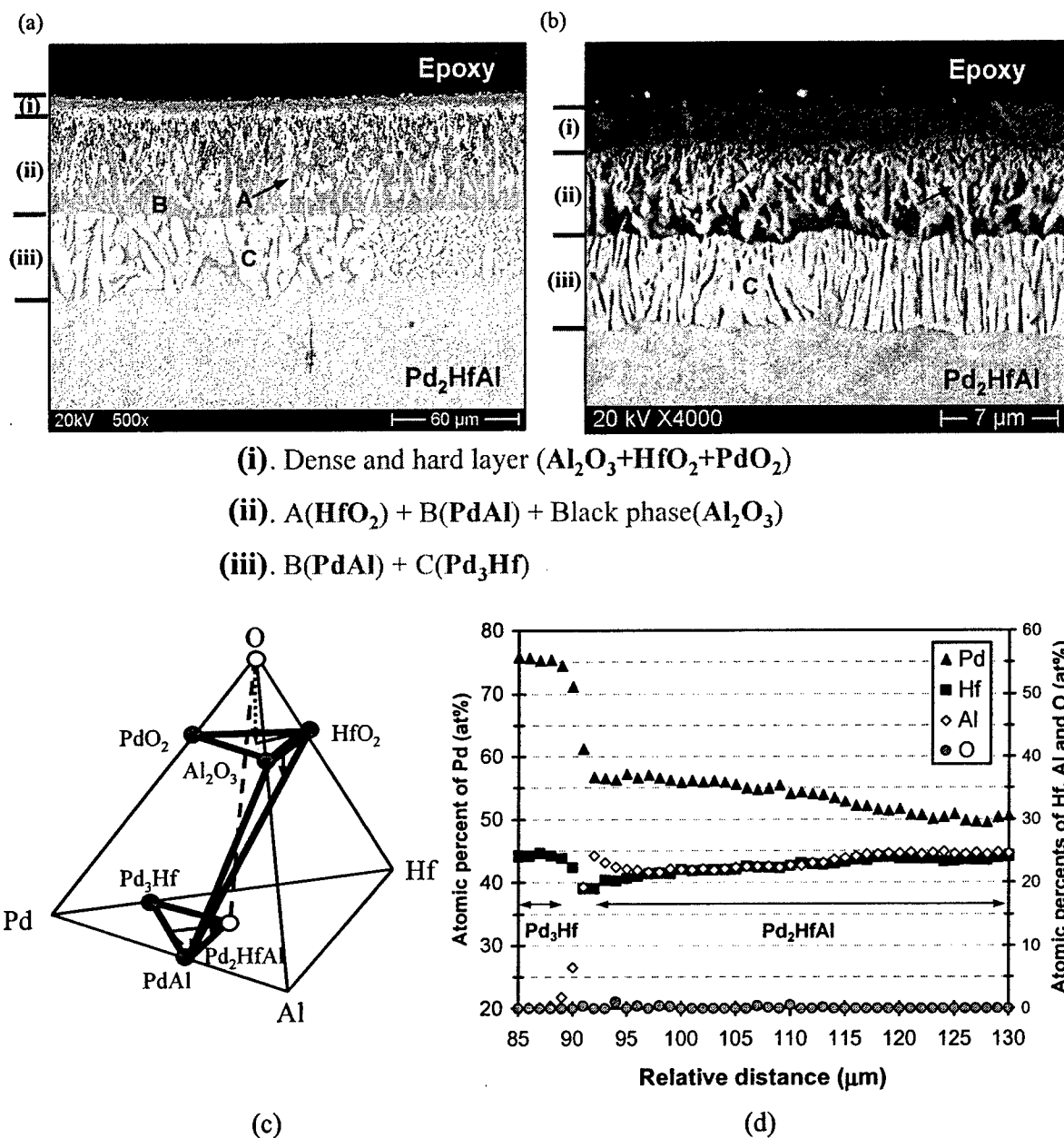


Figure 5. Cross section SEM-BSE images of the oxidized Pd_2HfAl alloys at (a) 1100 °C and (b) 1300 °C, showing three distinct layers: the (i) layer consists of Al_2O_3 , HfO_2 and PdO_2 , the (ii) layer consists of HfO_2 , PdAl and Al_2O_3 (black phase), and the (iii) layer consists of PdAl and Pd_3Hf . (c) Schematic oxidation reaction pathway and (d) concentration profiles of Pd, Hf, Al and O measured over Pd_3Hf - PdAl two-phase region and Pd_2HfAl region.

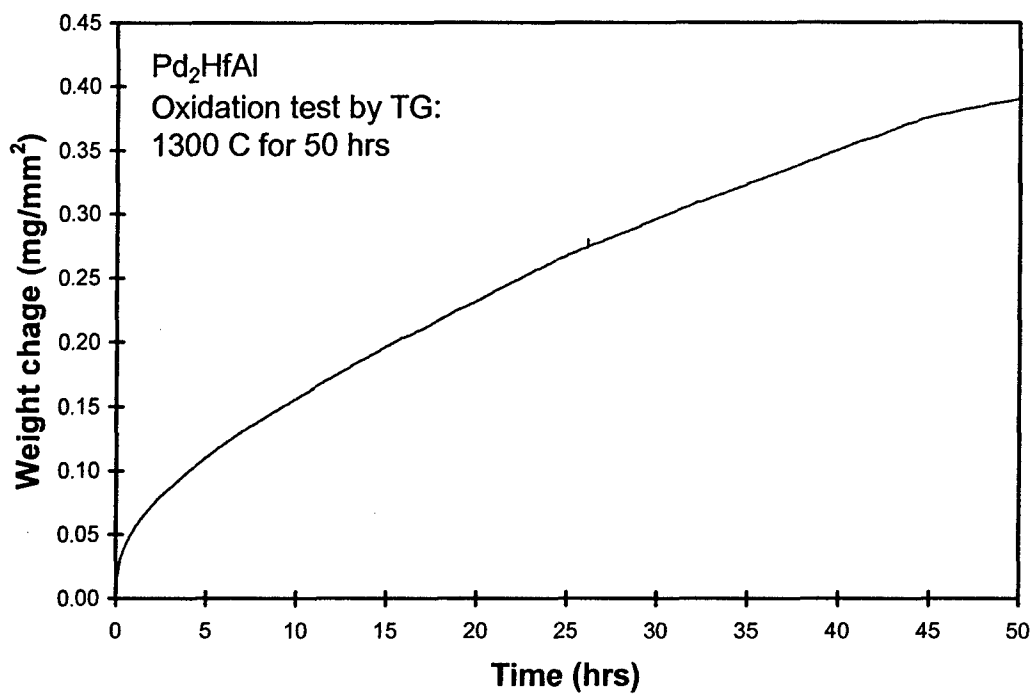


Figure 6. Plot of weight change of Pd_2HfAl versus oxidation time, obtained from oxidation experiment at 1300 °C by the TGA.

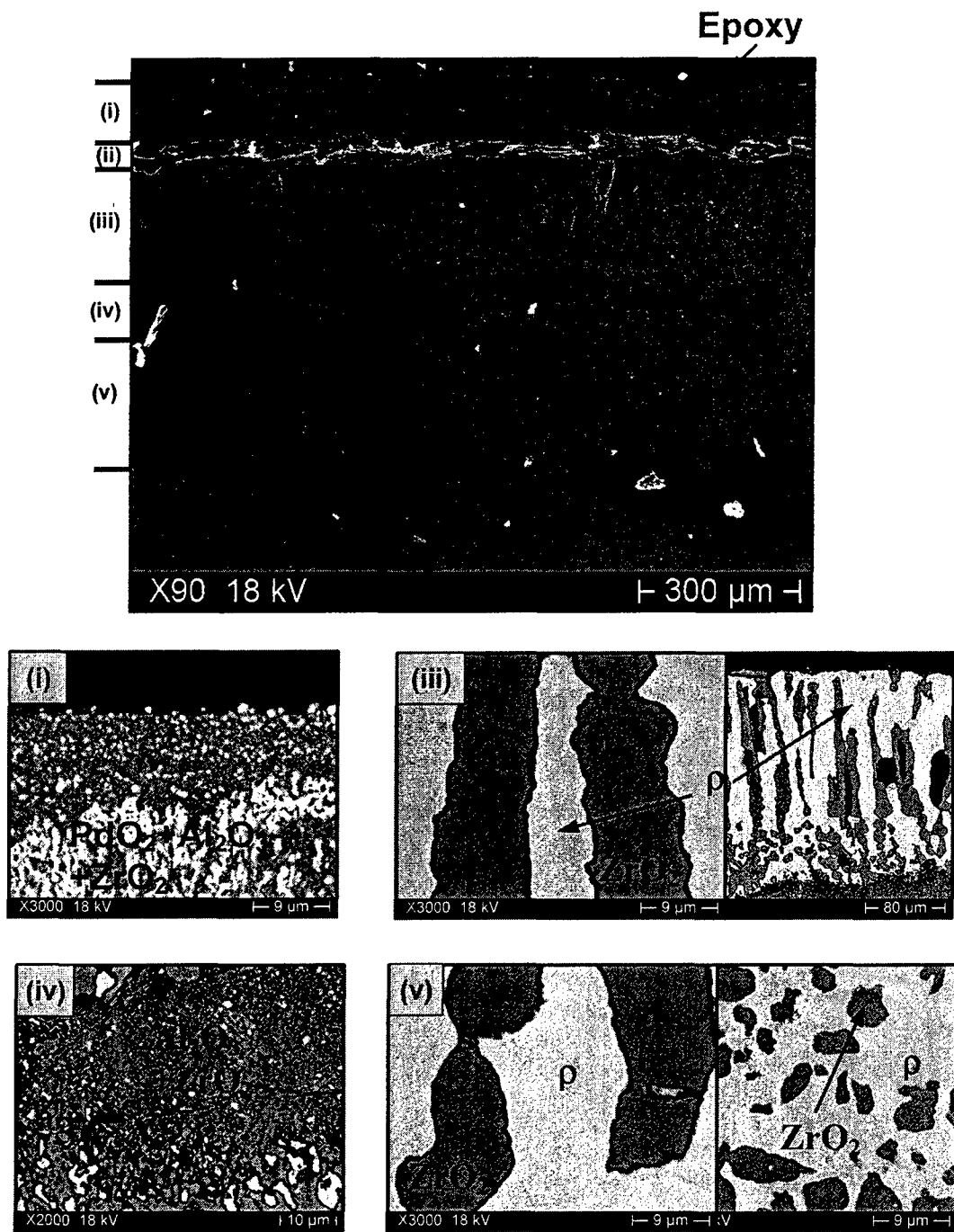


Figure 7. SEM-SE image of oxidized Pd_2ZrAl alloy at 1100 °C for 25 hours, showing five distinct layers: the (i) layer consists of PdO_2 , Al_2O_3 and ZrO_2 , the (ii) layer is a ZrO_2 oxide layer, the (iii) layer consists of ZrO_2 and ρ ($\text{Pd}_{75}\text{Al}_{25}$), the (iv) layer consists of Al_2O_3 and ZrO_2 , and the (v) layer consists of ZrO_2 and ρ .

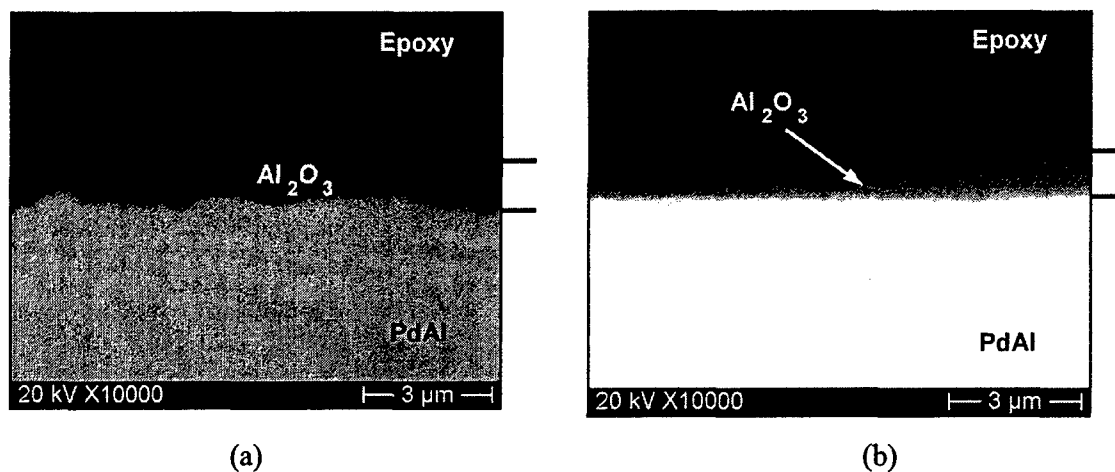


Figure 8. Cross section SEM-BSE images of the oxidized PdAl alloys at (a) 1100 °C and (b) 1300 °C for 25 hours.

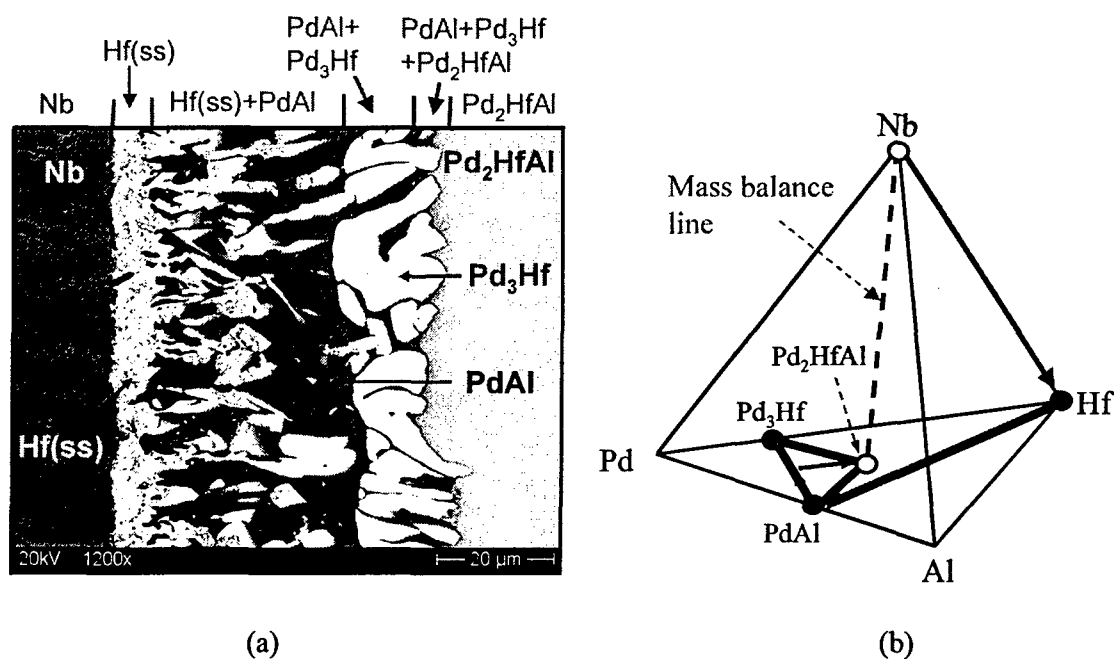


Figure 9. (a) SEM BSE image of the cross-section of the Nb/Pd₂HfAl diffusion couple annealed at 1100 °C for 50 hours and (b) schematic diffusion pathway.

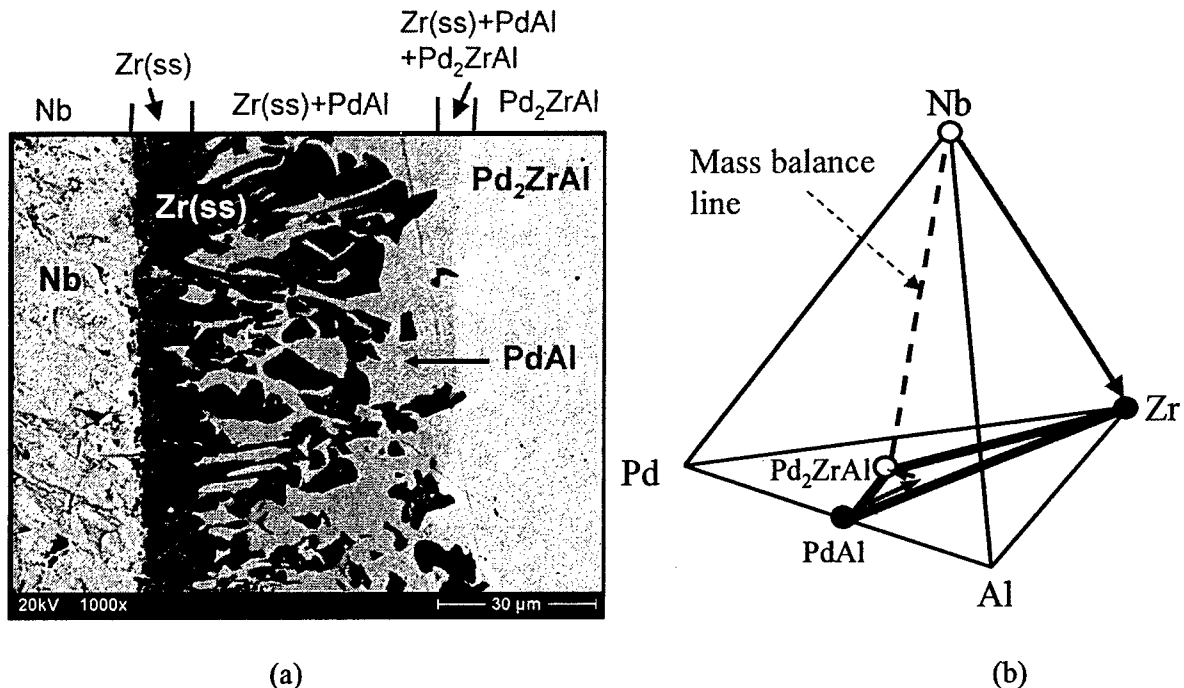


Figure 10. (a) SEM BSE image of the cross-section of the Nb/Pd₂ZrAl diffusion couple annealed at 1100 °C for 50 hours and (b) schematic diffusion pathway.

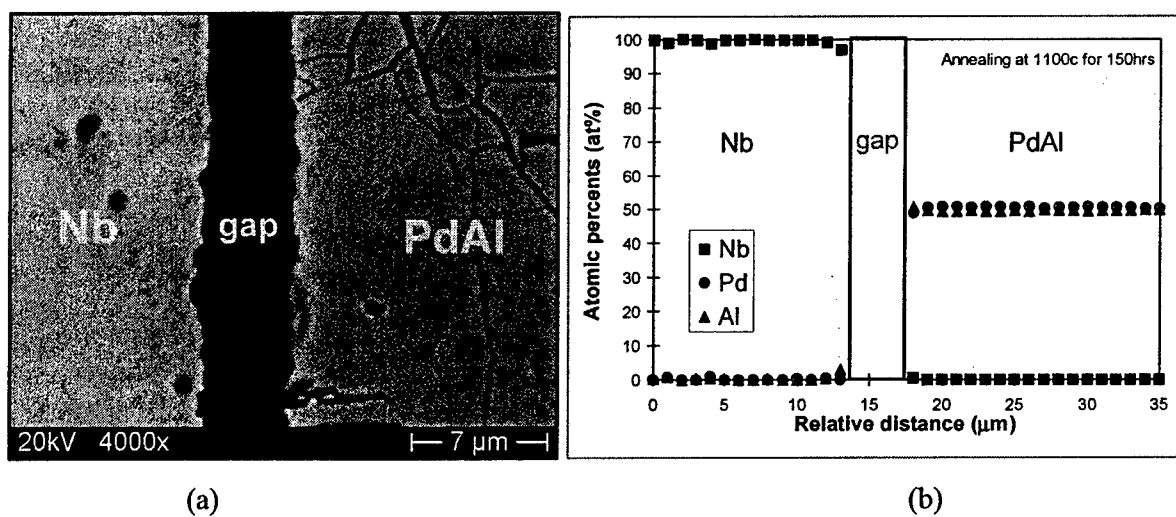


Figure 11. (a) SEM-BSE image of a cross section of the Nb/PdAl diffusion couple annealed at 1100 °C for 150 hours and (b) Nb, Pd and Al concentration profiles obtained from an EPMA line-scanning on the diffusion couple shown in (a).

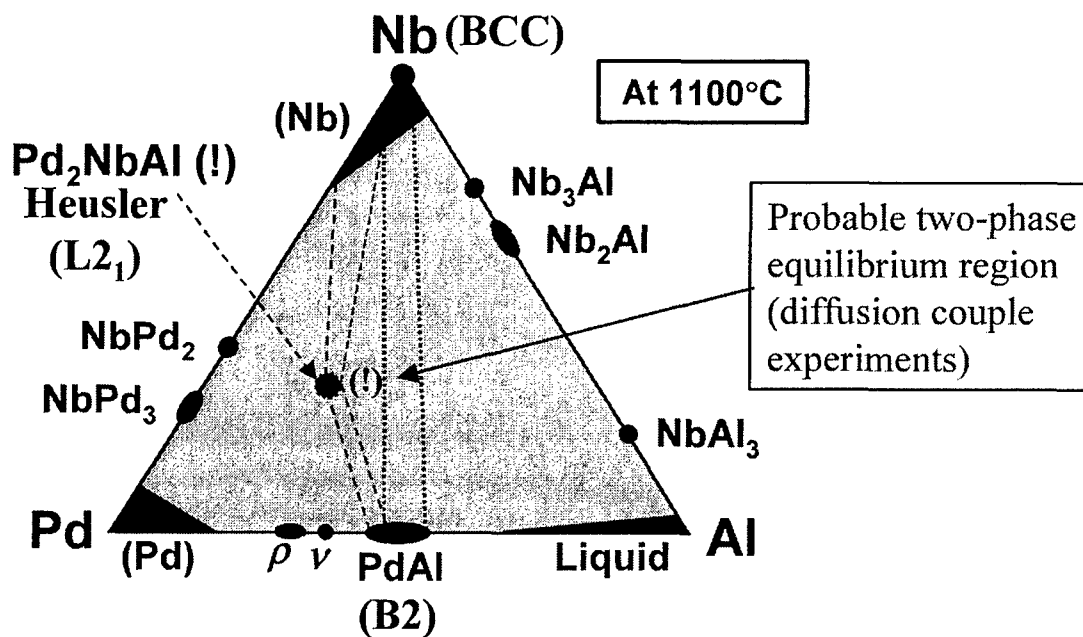


Figure 12. Schematic ternary phase diagram of the Pd-Nb-Al system showing probable three-phase equilibrium among bcc(Nb), B2(PdAl) and L2₁(Pd₂NbAl).

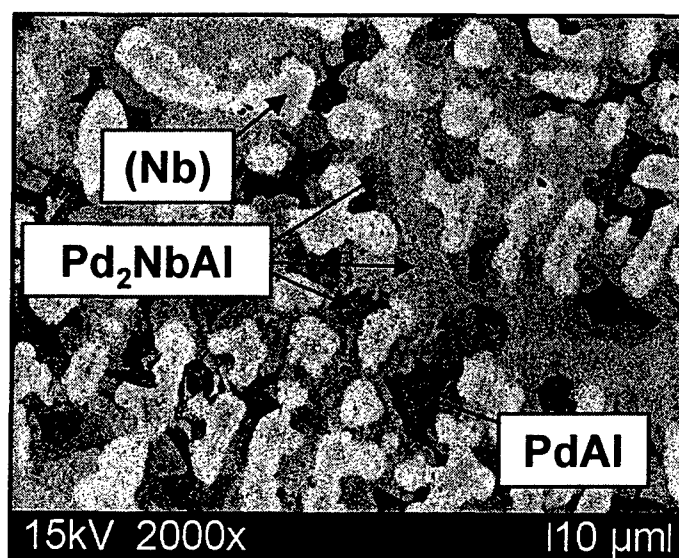


Figure 13. SEM-BSE image of as-cast Pd-25Nb-25Al (at% in unit) alloy showing co-existence of (Nb), PdAl and Pd₂NbAl.

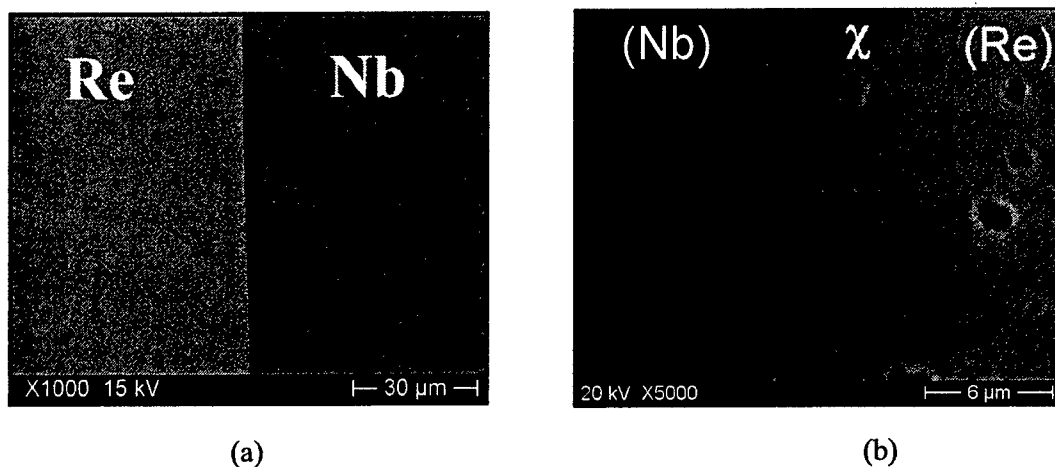


Figure 14. Cross section SEM-BSE image of the Nb/Re diffusion couple annealed at (a) 1100 °C for 200 hours and (b) 1600 °C for 100 hours.

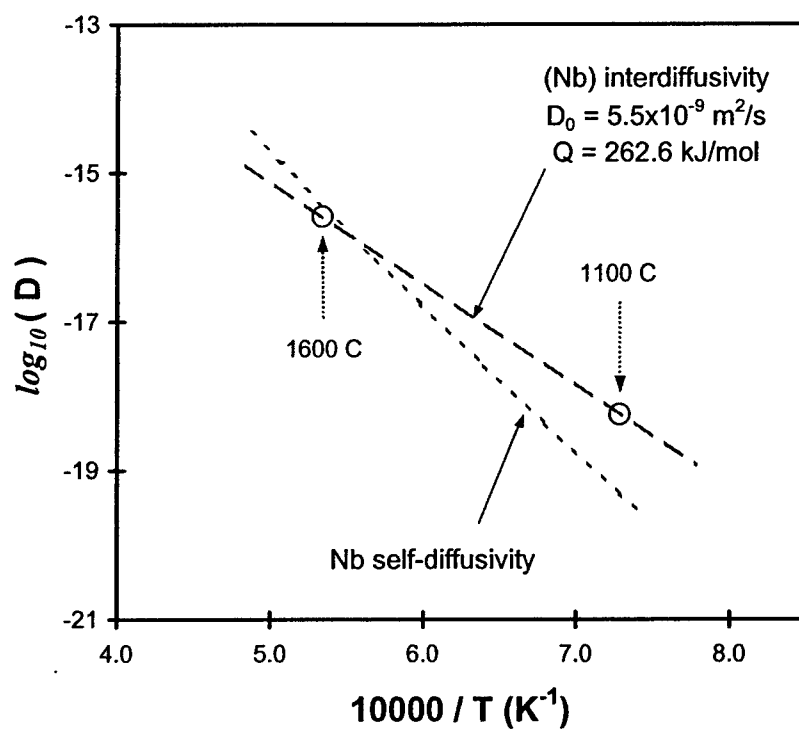
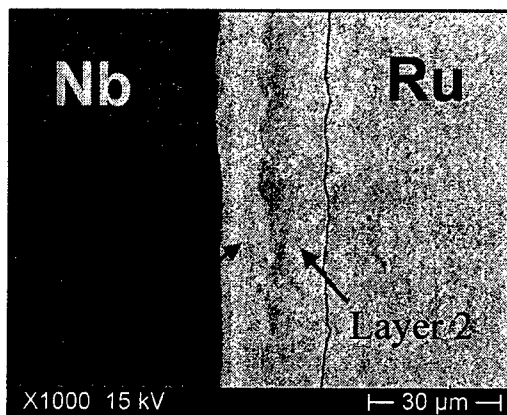
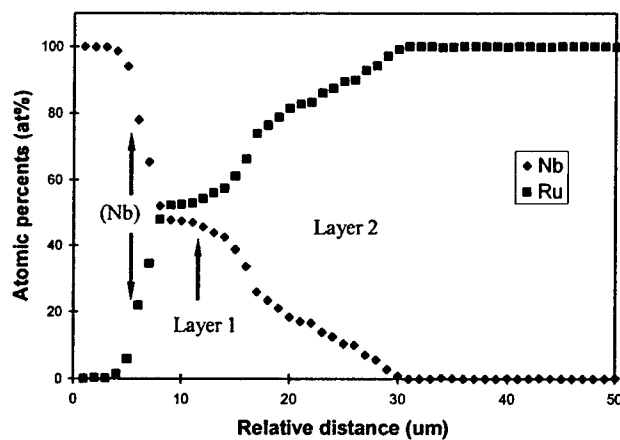


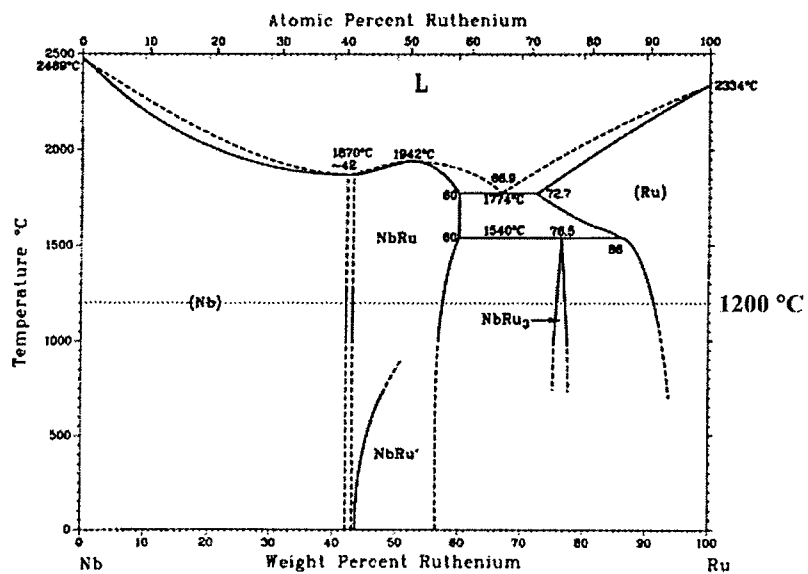
Figure 15. Plot of diffusion coefficients versus reciprocal temperature.



(a)



(b)



(c)

Figure 16. (a) SEM-BSE image of a cross section of the Nb/Ru diffusion couple annealed at 1200 °C for 200 hours, (b) Nb and Ru concentration profiles obtained from an EPMA line-scanning on the diffusion couple shown in (a), and (c) the Nb-Ru binary phase diagram in reference *asm96*.

REPORT DOCUMENTATION PAGE

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