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*G. R. Smolik
D. A. Petti
J. P. Sharpe
S. T. Schuetz*

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**OXIDATION AND VOLATILIZATION FROM
TANTALUM ALLOY T-222 DURING AIR EXPOSURE**

**Galen R. Smolik
David A. Petti
J. P. Sharpe
Stan T. Schuetz**

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**Idaho National Engineering and Environmental Laboratory
Idaho Falls, Idaho 83415**

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ABSTRACT

Tantalum alloys are one of the refractory metals with renewed consideration for high temperatures in fusion reactor applications. Tantalum alloys perform well in protective environments but oxidized readily in gases containing higher oxygen levels. In addition, the radioactive isotope Ta-182 would be produced in tantalum and could be a significant contributor to dose if mobilized. Other isotopes of importance are produced from tungsten and hafnium. Mobilization of activated products during an accident with air ingress is therefore a safety issue. In this study, we measured the extent of oxidation and mobilization from tantalum alloy T-222 oxidized in flowing air between 500 and 1200°C. This alloy nominally contains 10 wt% tungsten, 2.5 wt% hafnium and 0.01 wt% carbon. We found that the mobilization of Ta and Hf was closely linked to the occurrence of oxide spalling. These elements showed no migration from the test chamber. Some W was mobilized by volatilization as evidenced by transport from the chamber. Tungsten volatilization could occur primarily during initial stages of oxidation before an oxide scale forms and impedes the process. The mobilization of Ta and W are presented in terms of the mass flux ($\text{g}/\text{m}^2\text{-h}$) as a function of test temperature. These measurements along with specific designs, activation calculations, and accident scenarios provide information useful for dose calculations of future fusion devices.

SUMMARY

In this report we present the results of experimental measurements of the oxidation and mobilization of tantalum, tungsten and hafnium from the tantalum alloy T-222. Specimens of the alloy were exposed to flowing air between 500 and 1200°C. After exposure the extent of oxidation and the character of the oxide were determined by mass change, optical and electron microscopy, x-ray diffraction, and particle size analysis. The mobilization of the three elements mentioned above were determined by chemically removing the oxides from the test components, dissolving the oxides, and analyzing the solutions with inductively coupled plasma-atomic emission spectroscopy (ICP-AES).

The experimental results are presented in Section 3. Photographs of the specimens in Section 3.1 provide good indications of the amount of oxide spalled and powder formed at the various test conditions. The amounts and rates of oxidation as determined from mass change and alloy recession are discussed in Section 3.2 (and reported in Table 1). Illustrations of the oxide scales formed at the various temperatures as revealed by SEM are presented in Section 3.3. Powder was formed most abundantly at 700°C. We performed particles size analysis by various methods and have reported the results in Section 3.4. Metallographic cross sections were used to show the features and thicknesses of oxide scales and internal effects from oxidation in Section 3.5. Internal effects included dissolved oxygen that increased hardness and made the alloy brittle. Phase determinations from x-ray diffraction are reported in Section 3.6. Mass flux determinations ($\text{g}/\text{m}^2\text{-h}$) for tantalum, tungsten and hafnium at the various temperatures are presented in Section 3.7.

Experimental results are discussed in Section 4. We conclude that the mobilization of tantalum and hafnium is linked to the oxide spalling process because the oxides of these elements have very low volatility. Tungsten on the other hand has very volatile WO_3 polymeric species. We found that some tungsten was mobilized by volatilization. This was evident by increased mobilization with temperature and transport out of the reaction chamber. The mobilization of tantalum actually decreased with temperature which correlated with less particulate formation. We found little evidence of tantalum transport beyond the test chamber. This indicates that the oxide powder that spalled from the specimens has a low susceptibility to being transported. Upon comparing mobilization rates with a previously tested powder metallurgy (PM) alloy, containing 95 wt % tungsten and 0.1 wt % tantalum, we find that the mobilization of tungsten from the T-222 is actually less than that predicted based upon alloy content. This suggests that an oxide scale forms and impedes tungsten loss by volatilization. Mobilization rates of tantalum from the PM product, however, were higher than those from the T-222. This exemplifies the influence of oxide spalling and the importance of considering thermal cycling and other factors on the mobilization from tantalum alloys during accidents involving potential oxidation. Oxygen dissolution, embrittlement and impact upon mechanical properties are other issues to consider during off-normal conditions involving the oxidization of tantalum alloys.

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1. INTRODUCTION

Extensive effort was directed during the mid-1960's to develop tantalum alloys for use for space nuclear reactor systems.[1] Minor oxygen contamination in the alloys promoted corrosion in liquid lithium. Improvements were achieved by alloying small concentrations of hafnium to combine with the oxygen. Alloy T-222 represents one of these improved alloys. Tungsten and a small amount of carbon were added to improve high temperature creep strength. Such alloys have received renewed interest within the fusion community for use in liquid metal coolant systems. Potential mobilization of activated products during operation or accidents with air ingress is a safety concern. Tantalum alloys readily dissolve oxygen and oxidize above 500°C. Oxidation kinetics are complexly influenced by oxygen levels, oxygen dissolution, and the nucleation and growth of substoichiometric oxides of various oxidation states.[2,3,4] The solubility of oxygen in tantalum ranges from 2 to 4 atomic percent between 800 and 1200°C. After dissolving sufficient oxygen into the matrix heterogeneous nucleation of rod or plate-like metallic suboxides referred to as TaO_x , TaO_y , and TaO_z can occur. An oxide scale of β -phase Ta_2O_5 ultimately forms at high temperatures and oxygen levels. Oxidation kinetics progressively change from logarithmic, to parabolic, to linear modes with higher temperatures and oxygen pressures and longer times.[2,3] We would expect linear oxidation kinetics to readily occur at 600°C in air based upon the report by Kofstad.[2] Tantalum oxides have low vapor pressures. Volatilization has been reported to occur at low pressures, e.g., 10^{-5} torr O_2 and high temperatures (1600 to 1700°C). Vaporizing species at these conditions are TaO and TaO_2 .[2] Hafnium oxide is also very stable with negligible vapor pressure. Tungsten forms various volatile WO_3 polymeric species. We have measured volatilization from tungsten alloys.[5,6] The volatilization of WO_3 is enhanced by even small amounts of water vapor. The objectives of this study are to characterize the nature of the oxide scales formed on the T-222 alloy at various temperatures during the air exposures, to quantify mobilization of Ta, W, and Hf, and verify the mechanisms associated with the mobilization of each of these elements.

2. EXPERIMENTAL PLAN

Specimens were prepared from a 0.1-in. thick plate of Alloy T-222. The nominal dimensions of the specimens were 2.54 cm x 2.54 cm x 0.25 cm. They had a mass of about 30 grams and a surface area of about 15 cm². They were polished to a 600-grit finish and cleaned with ethanol. The specimens were tested in a 4 cm diameter quartz chamber heated in a 36 cm long muffle furnace. Air supplied from a cylinder was flowed through a column of Drierite and preheated to 400°C. The air was vented to the atmosphere at the ambient pressure at the INEEL of about 650 mm Hg. A flow rate of one liter per minute (STP) was maintained during all tests. This flow rate and the atmospheric pressure at the INEEL pressure produced gas velocities at specimen surfaces ranging from 0.04 to 0.08 meters per second over the temperature range of 500 to 1200°C. The gas then flowed through a 1.5-cm diameter quartz tube packed with quartz wool. The latter components extended from the furnace reaching near ambient temperatures at the outlet. The specific test temperatures and times are given in Table 1. The test durations were varied to provide longer exposures at lower temperatures. Mass changes of specimens were measured and oxides characterized. The oxidized specimens were examined and photographed. Cross sections of selected specimens were prepared and metallographically examined. Oxidized specimens and oxides were also examined using scanning electron microscopy, x-ray diffraction, and particle size analyses by image analyses and a Coulter particle size analyzer. Particles of oxides transferred to the quartz components were removed and dissolved with solutions of hydrofluoric and nitric acid. These solutions were chemically processed and analyzed by ICP-AES. Distributions of mobilized elements within the test system were thus determined.

Table 1. Test matrix and results from tantalum alloy T-222 tested in air.

Specimen	Temperature, °C	Time, h	Mass gain, g	Rate of oxygen pickup, g/h	Oxygen content, at%	Alloy loss per side, mm	Oxidation rate, mm/h
Ta1	500	81	0.0094	0.0001	0.4	0.07	9E-4
Ta2	600	55	1.9134	0.0348	41.9		
Ta3	700	55	0.5502	0.0100	19.2		
Ta4	800	48	3.9742	0.0828	60.0		
Ta5	900	8	1.7391	0.2174	39.6	0.27	3.4E-2
Ta6	1000	6	3.0939	0.5157	53.9		
Ta7	1100	4	2.1096	0.5274	44.3		
Ta8	1200	2	1.5479	0.7740	36.9	0.27	1.4E-1

(a) The nominal mass of the specimens is 30 g. Specimen mass was weighable to 0.0001 gram. Measurements of the metal core of specimens varied by ± 0.02 mm for the 500°C test and by ± 0.1 mm for the tests at 900 and 1200°C.

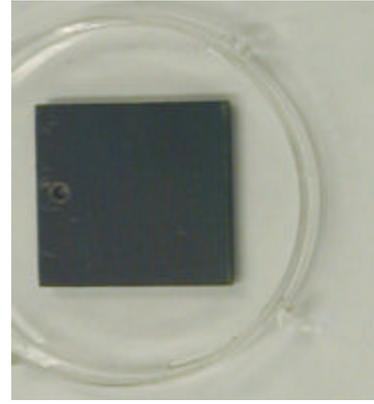
3. EXPERIMENTAL RESULTS

3.1 Visually Apparent Features of Oxidized Specimens

The extent of visually apparent oxidation and oxide spalling from specimens and in the test chambers provided insight into mobilization rates. Photographs of the specimens tested between 500 and 1200°C are shown in Figure 1. The specimen tested at 500°C did not appear significantly different than that of the initial untested specimens. It had a very thin dark oxide. Transition to an advanced stage of oxidation is indicated by a voluminous light yellow oxide forming at one edge. The specimen exposed at 600°C had developed a complete scale of yellow oxide. The flat sides and edges had spalled off as large pieces. There was also an oxide powder released from beneath the scale. Some particles of this oxide were apparent in the test chamber. The yellow oxide that developed on the specimen at 700°C had turned almost completely into powder. Some of this powder was present in the reaction chamber. The specimen tested for 48 hours at 800°C had developed about a 2.5-mm thick oxide scale. The sides and edges had spalled off revealing a thin dark gray core. Some fine yellowish powder was released along with larger flat grayish particles from the zone near the oxide to metal interface. Less of the fine yellowish powder was released when the oxide scales spalled from specimens exposed at 900°C and 1000°C. The pieces of scale appeared to be sintered into a harder oxide. Oxide scales remained mostly intact after the shorter exposures of 4 and 2 hours at 1100°C and 1200°C, respectively. The oxide on the flat surfaces of the specimens appeared partially transparent. This suggests that the oxide is likely tantalum pentoxide, Ta₂O₅. The CRC Handbook of Chemistry and Physics shows this oxide as being a colorless crystalline phase.[7] The sample appeared gray on the flat surfaces due to the underlying oxide at the alloy interface. The translucent nature of the oxide suggests that it was somewhat dense and adherent. A more voluminous yellow oxide formed at the cusp-shaped regions of higher oxidation at the corners of the specimens. This oxide was more typical of the oxides formed at lower temperatures. There were only minor amounts of particles evident in the quartz test chambers. A black deposit formed on the front end of the quartz wool during the 1200°C test. This was the only test to show such visually apparent deposit downstream from the specimen.

3.2 Measurements of the Extent of Oxidation

The amount of oxidation by the various specimens was obtained from mass change and change in the thickness of the metal core obtained from metallography. Oxygen pickup calculated from sample mass change is given in Table 1. These values have been used to calculate average rates of oxygen pickup (g/h) and the overall final oxygen content (at%) of the specimens. The rates of oxygen pickup increase progressively with temperature except for the test at 700°C. The oxygen pickup of 0.5502 g, or rate of 0.0100 g/h, at 700°C was less than one-third of that measured at 600°C. It is possible that some of the fine powdery oxide from the 700°C test could have been lost during handling. ICP-AES, however, showed that only about 1 mg of tantalum plus tungsten was recovered from the test chamber for that test. We do not believe that up to one gram of oxygen (or about 5 grams of oxide) could have been lost during handling. Rather, the difference between the 600 and 700°C test seems to reflect temperature trends of oxidation rates observed for Ta and Nb by other studies.[2,3,8] The transformation of metallic suboxide phases to pentoxides has been suggested as a possible reason for this behavior.



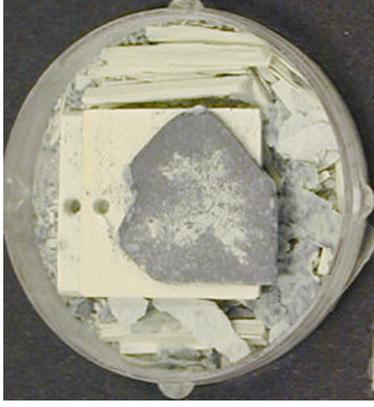
(a) 500°C/81 hours



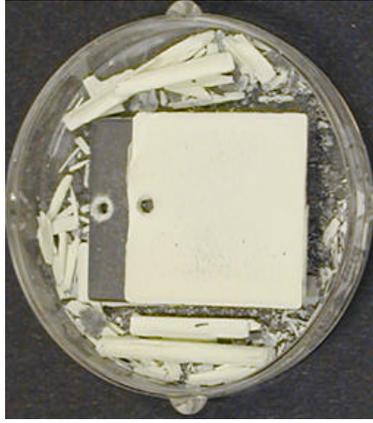
(b) 600°C/55 hours



(c) 700°C/55 hours



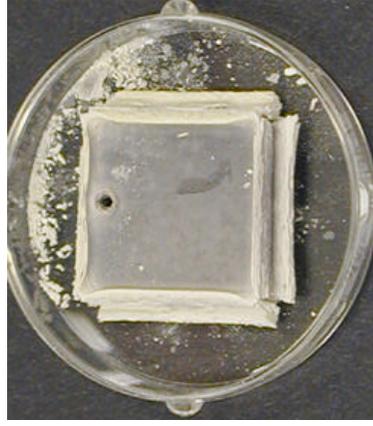
(d) 800°C/48 hours



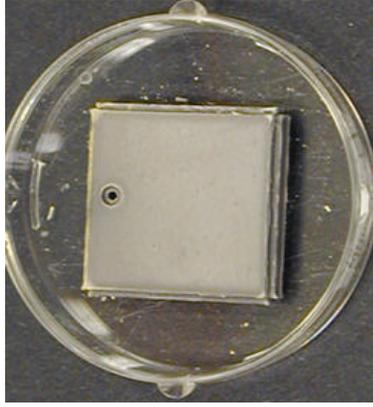
(e) 900°C/8 hours



(f) 1000°C/6 hours



(g) 1100°C/4 hours



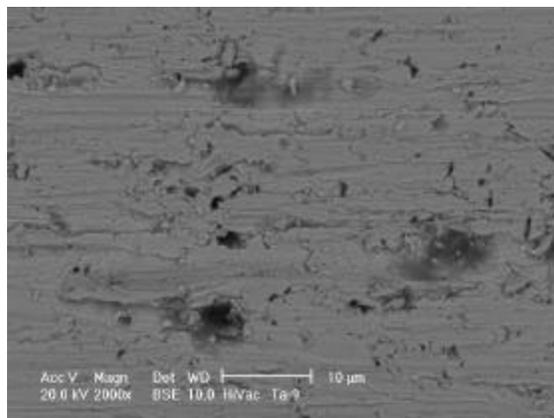
(h) 1200°C/2 hours

Figure 1. Appearance of tantalum Alloy T-222 specimens tested in flowing air at various times and temperatures.

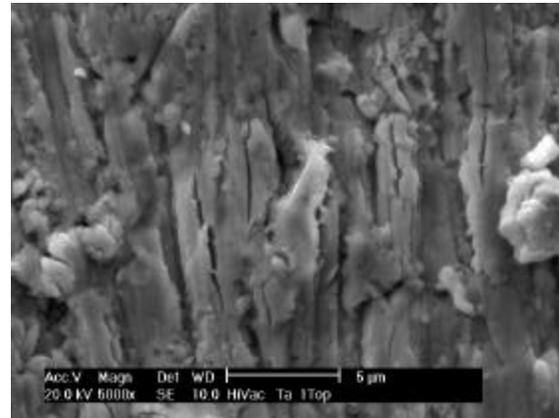
Data in Table 1 show that oxygen in the thin layer from the 500°C test contributes less than 0.5 at% to the specimen. The specimen tested at 800°C was the most completely oxidized. It attained an oxygen content of about 60 at%. Since some of the specimen core had still not been reacted, it is likely that much of the oxide had been converted to Ta₂O₅, which contains 71 at% oxygen. Alloy recession rates based on metallographic observations are shown in Table 1 for the specimens tested at 500, 900 and 1200°C.

3.3 SEM Depiction of Oxide Scales

SEM displayed the morphologies of oxides formed on the specimens. Figure 2 shows a comparison between surfaces of an untested as-polished specimen and the one tested at 500°C. The untested specimen in Figure 2(a) shows patterns of elongated smeared material formed during the polishing process. Such smearing is indicative of ductile material. Harder carbide inclusions stand out in relief. The SEM image of the specimen tested at 500°C in Figure 2(b) shows microcracks in the laminated oxides formed along the preexisting polishing marks. SEM showed progressive changes in oxide features with increasing temperature. A thicker more fibrous type of oxide formed at 600°C as shown in Figure 3(a). Oxide formed at 900°C shown in Figure 3(b) was shaped more like platelets. Some small rounded nuclei were beginning to form on the oxide platelets. Figure 3(c) and Figure 3(d) show the oxides formed at 1100 and 1200°C, respectively. The oxides are composed of small rounded crystals 0.2 to 1 μm in diameter. These small crystals appear to be bonded together. The oxides are relatively free of cracks and voids. These features may provide some impediment to the mobilization of tungsten as we discuss later in the interpretation of ICP measurements.



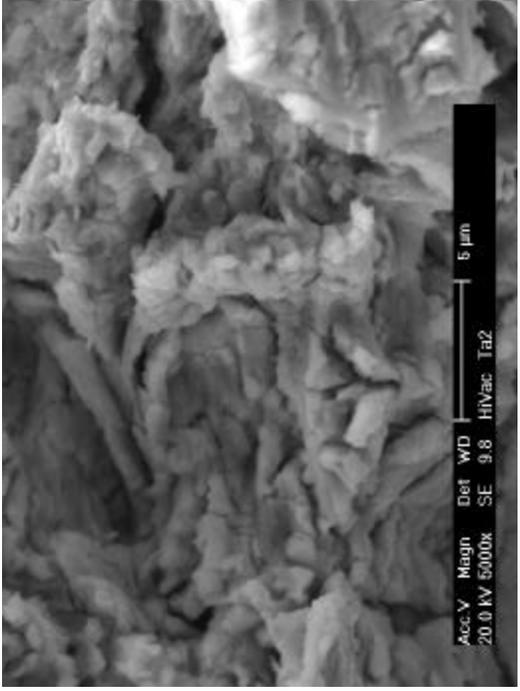
(a) As-polished specimen



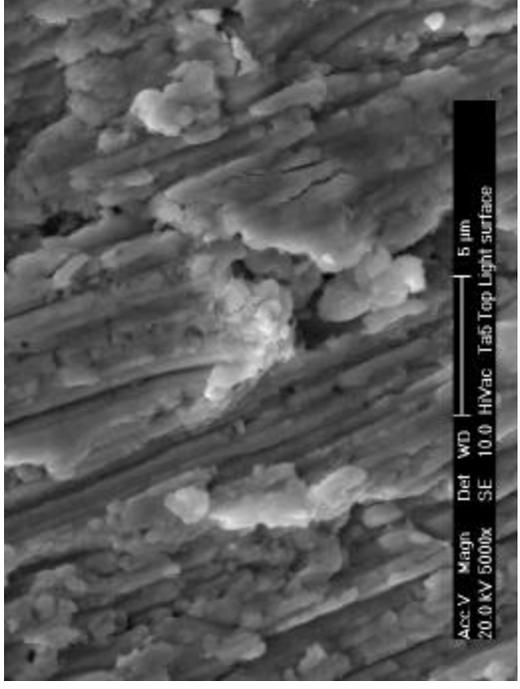
(b) Oxide on 500°C specimen

Figure 2. SEM images of the surfaces of T-222 specimens.

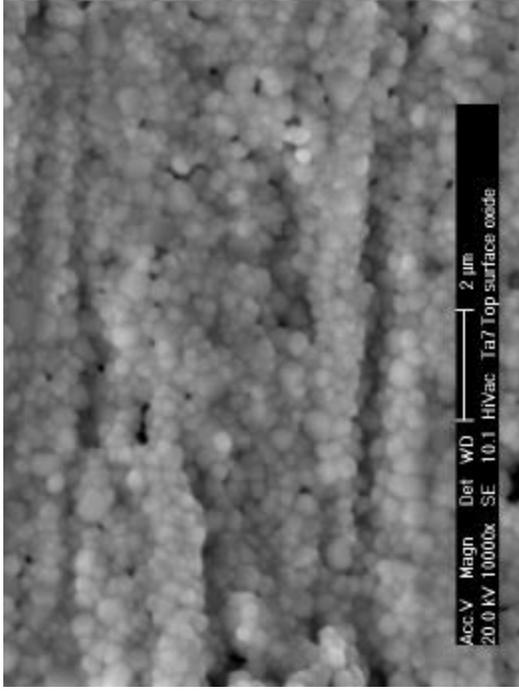
The sides of the specimen tested at 900°C spalled off as large pieces. The bottom surface of the oxide and the opposing surface from the specimen core are shown in Figure 4. The surfaces are similar. Both surfaces were relatively smooth and had very little texture or fibrous tearing. These are features indicative of a brittle type failure. The material shown may be either the oxide scale or the alloy along the interface that has dissolved a high amount of oxygen. We believe the spalling occurred during cooldown since the features are so sharp and distinct. This appears to be the source of the larger, flat, grayish particles observed in the powders.



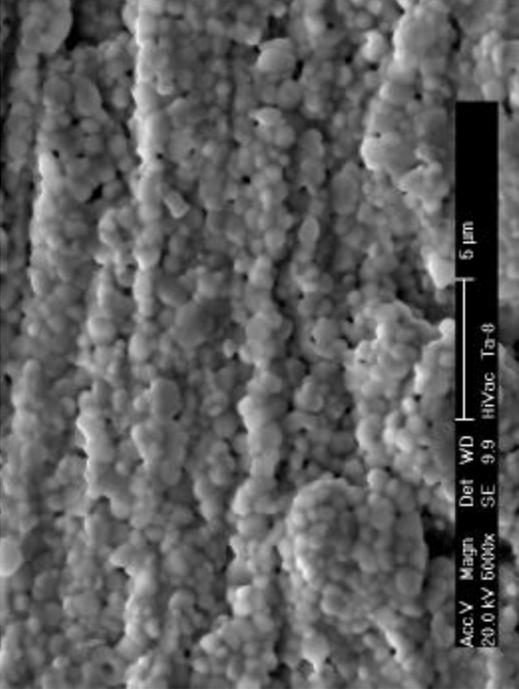
(a) 600°C/55 hours



(b) 900°C/8 hours

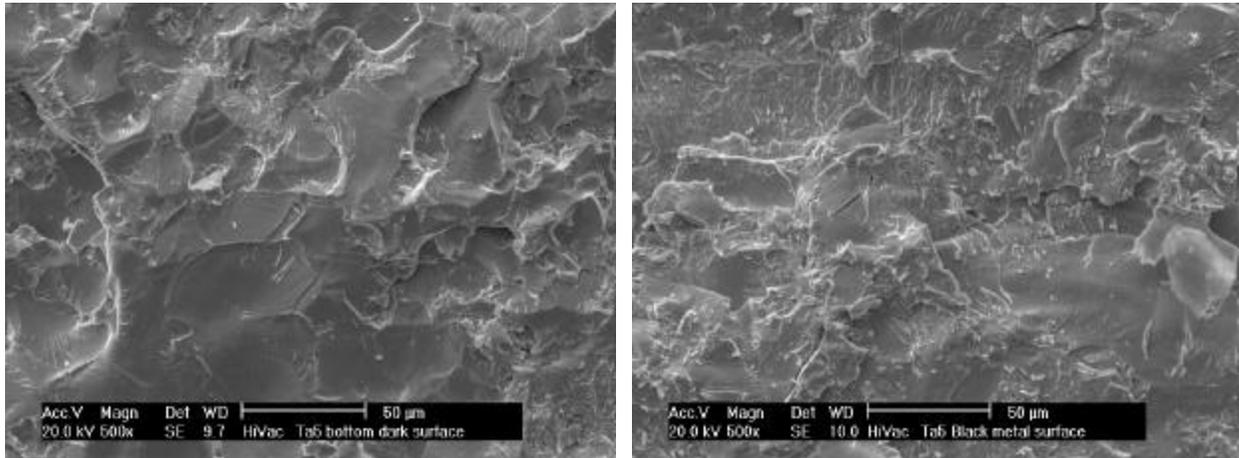


(c) 1100°C/4 hours



(d) 1200°C/2 hours

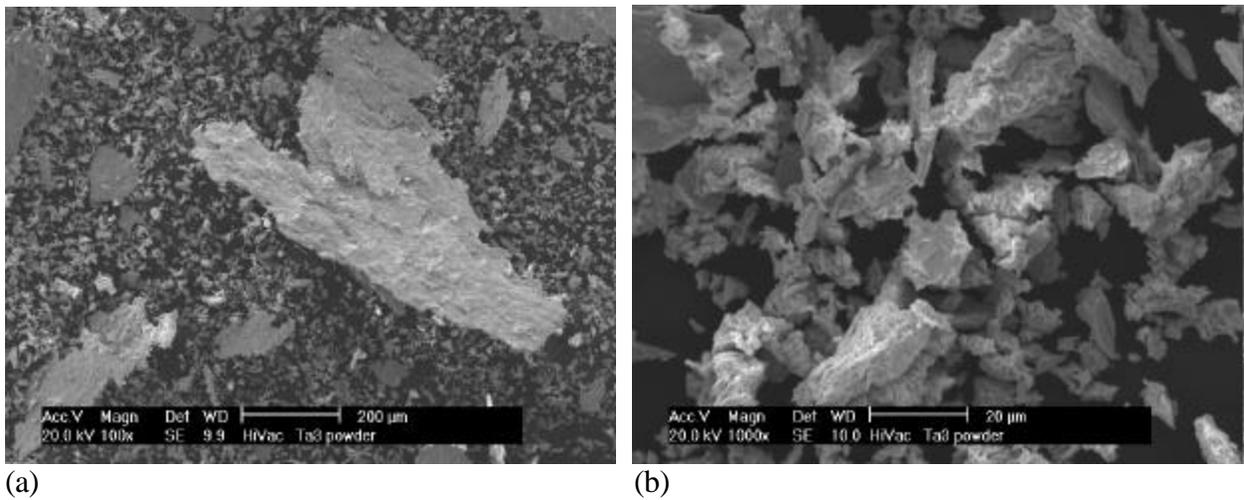
Figure 3. Outer surfaces of oxides formed on the tantalum T-222 specimens at various temperatures.



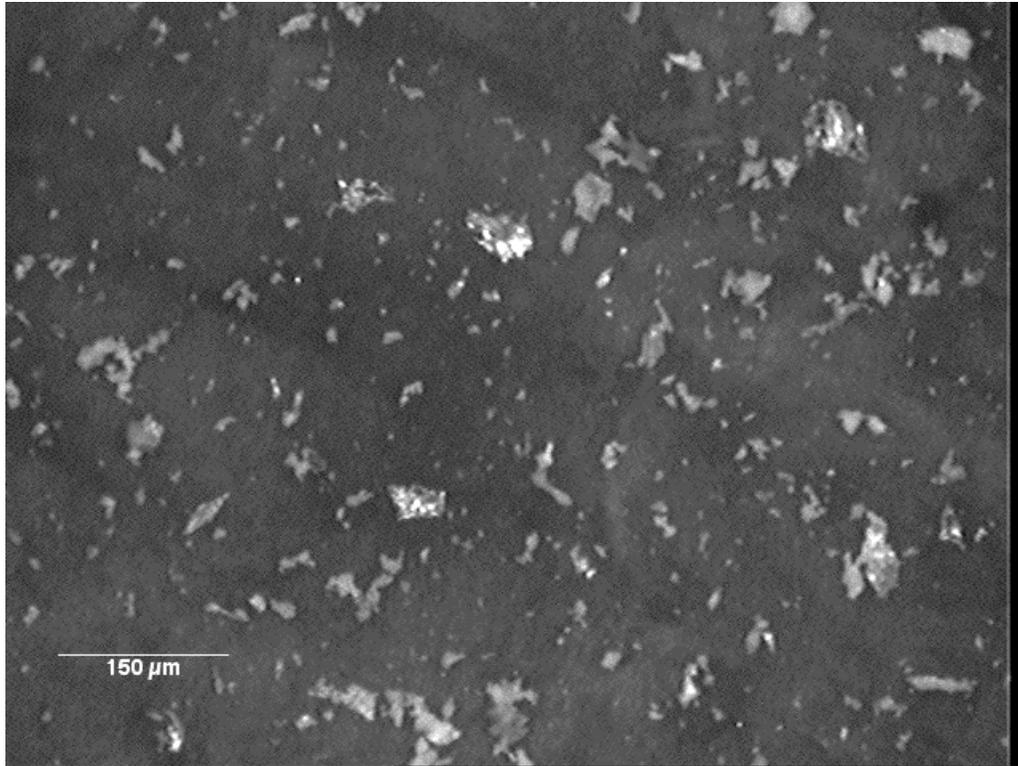
(a) (b)
Figure 4. Surfaces from specimen tested 900°C showing brittle type fracture. (a) Bottom surface of spalled oxide. (b) Surface of the remnant specimen core.

3.4 Particle Size Analysis of Oxide Powder

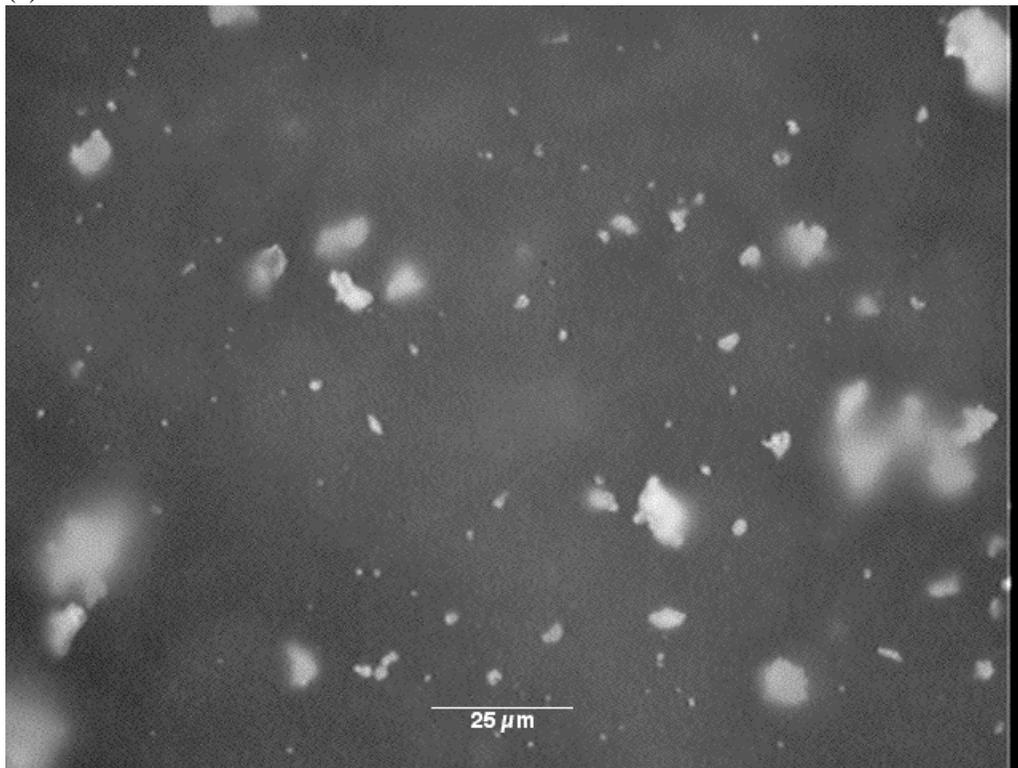
We selected the powder from the 700°C test to characterize due to the large amount of powder that formed. The SEM images in Figure 5 shows that there are different types of particles. The flat larger particles in Figure 5(a) are the gray particles we believe spalled from near the oxide to metal interface. The smaller irregular shaped particles in Figure 5(b) are those we believe were released from the voluminous outer yellow oxide. The latter type of particles was likely released during the oxidation process at the test temperature. We determined a count-based size distribution using optical microscopy and image analyses. Photographs were obtained from powder dispersed on a slide. Examples of the type of photographs used for the images are shown in Figure 6.



(a) (b)
Figure 5. SEM images of particulate formed from specimen tested at 700°C. (a) Illustration of various types of particles. (b) Irregular shape revealed for the smaller particles.



(a)



(b)

Figure 6. Examples of photographs used for image analyses. (a) Magnification of 100X. (b) Magnification of 500X.

Data obtained by counting the particles from the photographs are shown in Figure 7 and Table 2. Deviations from a straight line on the log-probability plot indicate the sampled population is not best described by a lognormal distribution. The data summary in Table 2 shows a geometric median diameter (GMD) of 1.95 μm and a geometric standard deviation (GSD) of 2.92. Parameters for a lognormal fit are given in Table 2 and compared with the counted-based data in Figure 8. Particles size distribution based on volume, or mass, were determined with a laser diffraction method with a Counter particle size analyzer. These measurements are compared with the conversion from the count-based fit assuming the Hatch-Choate relationship in Figure 9. The poor correlation shows the influence of the nonspherical, flakelike nature of the particles. The plot of the data in Figure 9 displays shapes characteristic of bimodal distributions. This most likely represents the smaller yellowish particles and the larger flat grayish particles we observed in the powder and as shown by SEM in the Figure 5.

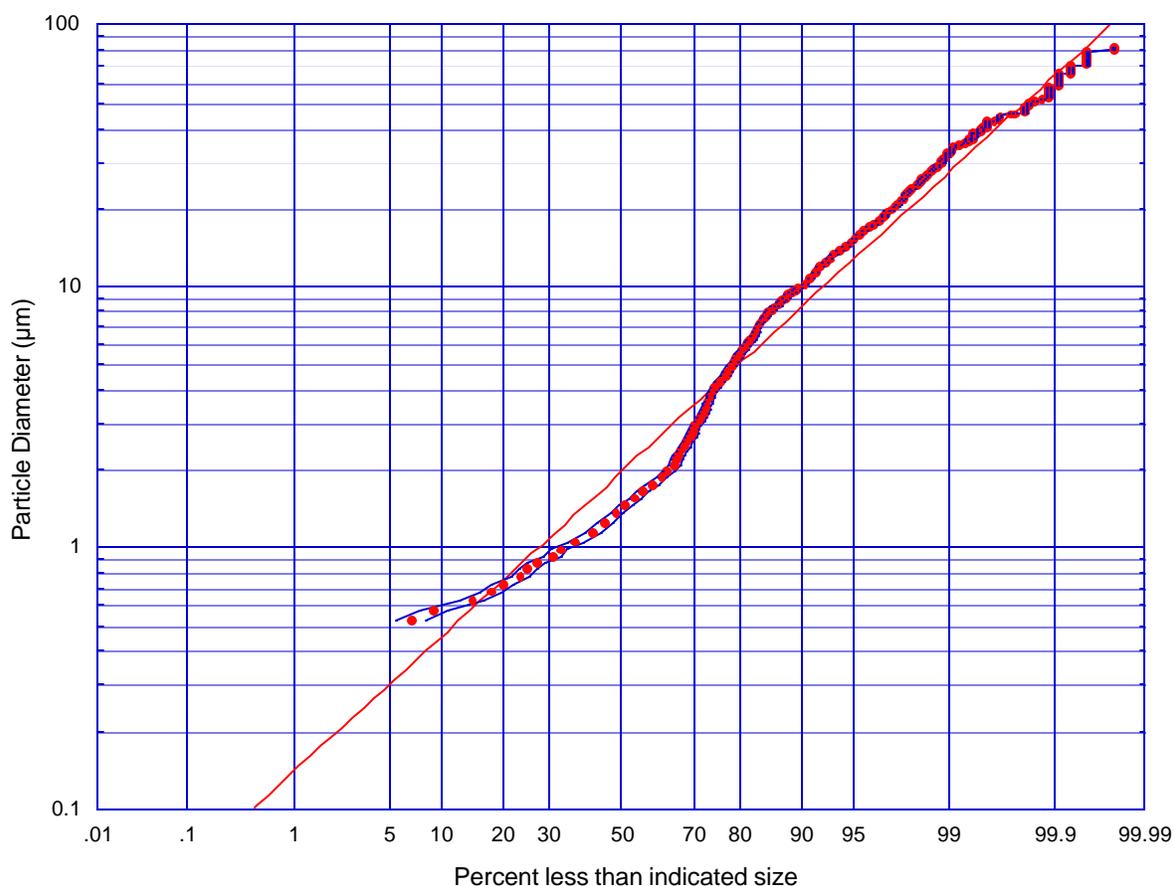


Figure 7. Log-probability plot of particles from the 700°C tests based upon a count-based analysis using optical photographs.

Table 2. Summary of count based particle size analysis.

magnification, number of images	sampled area ¹ (%)	counted size range (μm)	total number of particles (not scaled)
100x, 6	0.224	$8.00 < d_p$	1063
200x, 6	0.0558	$4.50 < d_p < 8.00$	641
500x, 6	0.00893	$2.10 < d_p < 4.50$	283
1000x, 6	0.00217	$0.00 < d_p < 2.10$	346
Totals:	1.18%	-	2,333
data moments ² :		log-normal fit moments ² :	
GMD (μm):	1.95	GMD (μm):	1.96
GSD:	2.92	GSD:	3.10
lower 95% (μm):	0.23	R ² :	0.98626
upper 95% (μm):	16.7	lower 95% (μm):	0.20
		upper 95% (μm):	18.8

1. based on overall filter area of $1.59 \times 10^9 \mu\text{m}^2$
2. GMD = geometric median diameter, GSD = geometric standard deviation, R = linear correlation coefficient

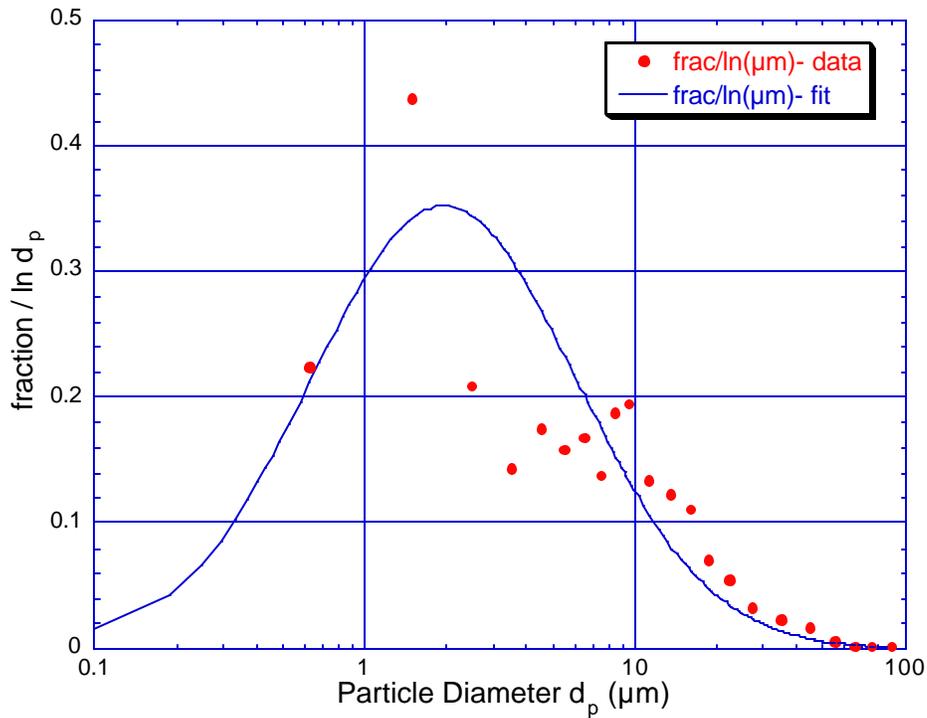


Figure 8. Equivalent count-based particle size distribution determined from data shown in Figure 7 and Table 2.

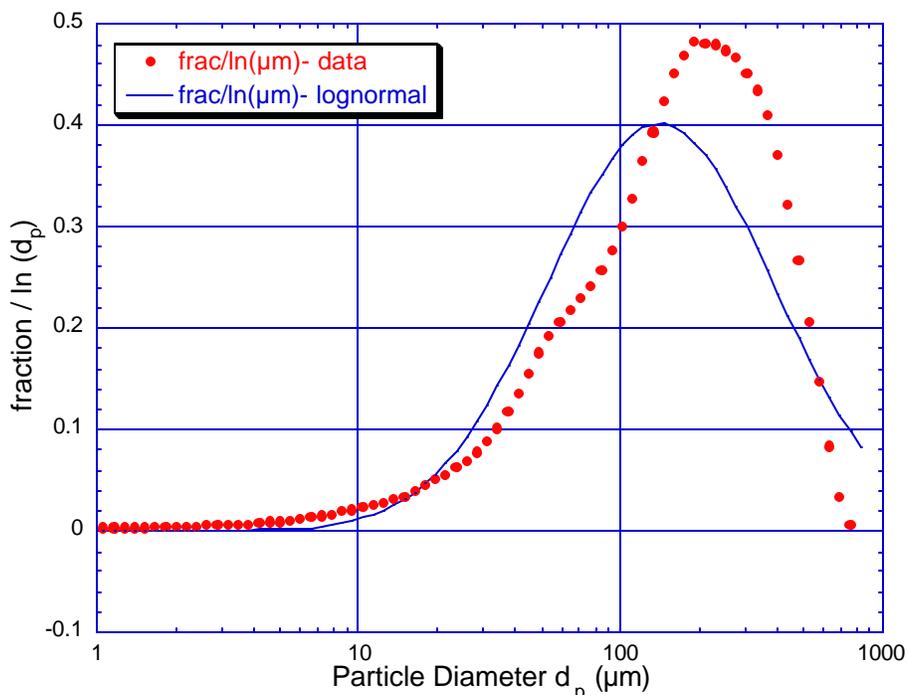


Figure 9. Data from volume- (or mass-) based size distribution for 700°C powder compared with a calculated lognormal distribution using Hatch-Choate relationship.

3.5 Internal Oxidation Revealed by Cross Sections

Besides revealing oxide scales and oxide to metal interfaces, metallographic cross sections showed evidence of an interstitial gas diffusion into the alloy. An example of the microstructure observed near the surface of the specimen tested at 500°C is shown in Figure 10. The tantalum carbides used for high temperature strength were not apparent in a 0.05-mm thick zone near the surface. Similar zones 0.2 mm thick were observed in the specimens tested at 900 and 1200°C. The high adsorption, diffusion and solubility of oxygen, and also nitrogen, into tantalum are well known.[2] The dissolution of these gases would strain and distort the lattice structure. This may enable carbides to be dissolved. It is possible that the precipitates were less apparent due to the polishing process on a hardened surface layer. The carbide inclusions, however, were not apparent with backscattered electrons with the SEM. This indicates that the regions near the surface are more chemically homogeneous than at interior locations where carbides exist. The diffusion of the interstitial atomic species into the alloy was illustrated by microhardness measurements. Diamond pyramid hardness profiles were made using a 50-gram weight. The hardness profiles for the specimens tested at 500, 900 and 1200°C are shown in Figure 11. All specimens showed increased hardness due to an apparent inward diffusion of some interstitial, presumably oxygen. The measurements in the zone within 0.05 mm of the surface void of carbides were significantly higher. The depth of diffusion extended to about 0.2 mm in the 500°C specimen. Microhardness readings then remained constant from there to the center of the specimen. Microhardness continuously decreased from the surface to the center of the specimen for the 900°C test. The values at the center approached those of the base material at 500°C. The

highest near-surface hardness was observed in the specimen tested at 1200°C. The hardness then decreased to nearly constant values at a depth of 0.5 to 0.75 mm. This showed that oxygen had not diffused completely to the center of the specimen during the two-hour exposure. The hardness within the core, however, was higher than that of the other two specimens. This suggests some change or restructuring of the carbides during the 1200°C exposure.

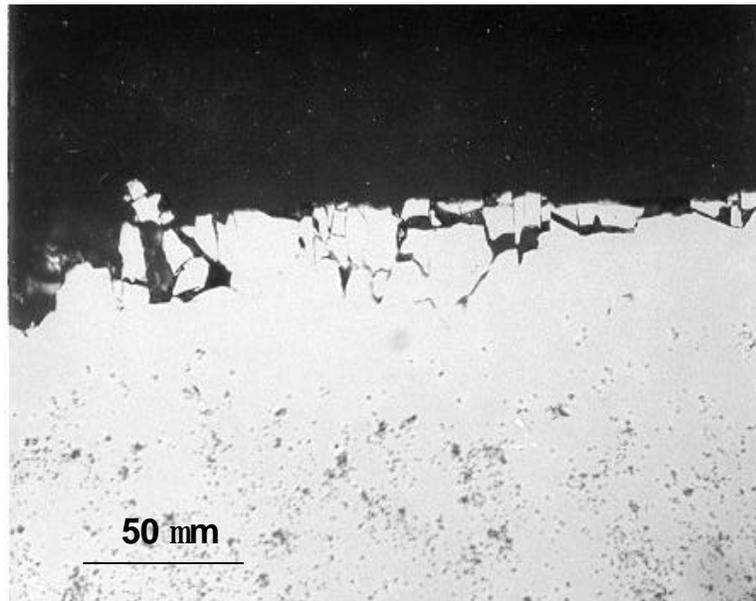


Figure 10. Zone near the surface of the T-222 specimen tested at 500°C that does not show the tantalum carbide precipitates.

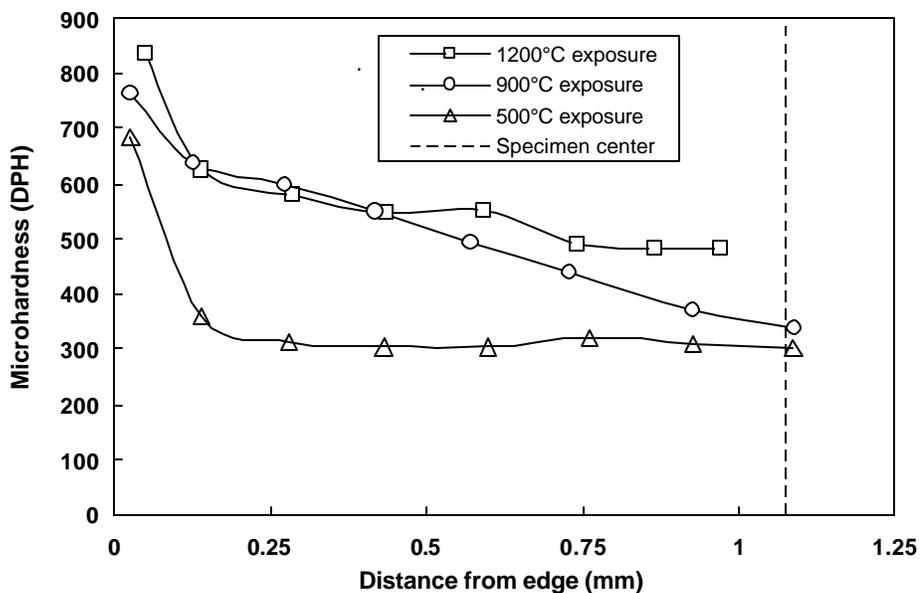
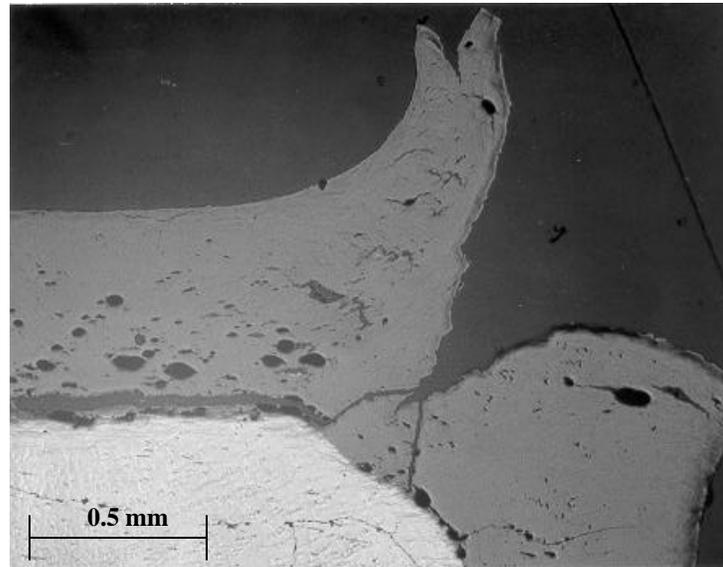
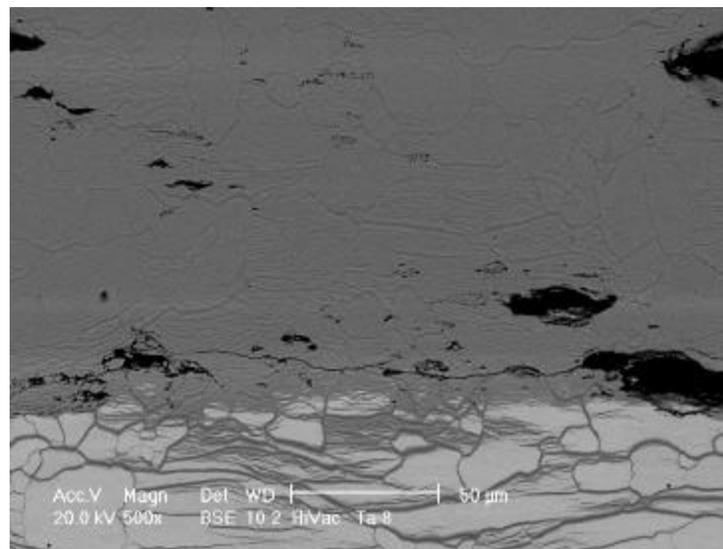


Figure 11. Microhardness profiles of specimens tested at 500, 900 and 1200°C showing the effect of oxygen diffusion into the specimen.

A consequence of dissolved oxygen is increased brittleness. Cracks exist within the hardened near-surface zone of the specimen in Figure 10. They are also apparent in the illustrations of the oxide scale and oxide to metal interface of the specimen tested at 1200°C in Figure 12. The cracks in Figure 12(a) extend deep into the metal substrate showing that the metal has become embrittled. The micrograph in Figure 12(a) shows that separation has occurred within the scale. The scale was still intact at other locations as shown in Figure 12(b). Thus the scale did not spall off during cooldown. Figure 12(a) shows some other features of the oxide scale. These are the enhanced oxidation resulting in the cusp shape at the corners, and porosity and cracking in the oxide scale due to the volumetric expansion from the oxide formation.



(a)



(b)

Figure 12. Cross section of oxide scale formed on T-222 specimen tested at 1200°C. (a) Optical micrograph at 50X. (b) SEM of oxide to alloy interface at 500X magnification.

3.6 Phase Identification by XRD

We performed x-ray diffraction (XRD) analyses of oxides from the 500, 700, 900, and 1200°C tests. The oxide scale was intact on the specimens tested at 500 and 1200°C. Oxide powders were used for the analyses for the 700 and 900°C exposures. A pattern expressed for a solid solution, Ta(O), formed from tantalum heated in air and oxygen between 470 and 520°C provided the best fit for the 500°C test.[9] There were some unidentifiable peaks that could result from lattice distortion due to dissolved oxygen. They did not match patterns for metallic tantalum phases. Diffraction peaks became more distinct with higher temperatures. Other oxide patterns, e.g., Ta₂O₅, and Ta_{14.8}W_{1.20}O_{40.6} provided fair fit to the peaks. The pentoxide pattern fit the spectrum from the 1200°C test quite well.

3.7 Mobilization Rates Obtained from ICP-AES Measurements

We present a summary of the mass flux determinations for tantalum, tungsten and hafnium from the ICP-AES measurements in Table 3. Detailed calculations for the mass flux are given in Appendix A. Although the reported flux values are derived from test chamber, the collection tube, and filter measurements, tungsten was the only element found in significant quantities beyond the test chamber. Hafnium was only detected within the test chamber at 600 and 700°C. These were the only measurements for this element that were above the Engineering Maximum Flux Value (EMFV) as previously defined.[10] Tantalum measurements were well above detection limits except for those for the 1100 and 1200°C tests. The measurements were only 20 to 40 percent above the detection limit at these higher temperatures. The mass flux determinations for tungsten were always greater than ten times the uncertainty.

Table 3. Mass flux determinations from ICP-AES measurements.

Temperature, °C	Time, h	Hf (g/m ² -h)	Ta (g/m ² -h)	W (g/m ² -h)
500	81	2.38E-5	1.12E-3	1.89E-4
600	55	2.48E-4	3.12E-2	2.64E-3
700	55	7.13E-5	1.30E-2	4.10E-4
800	48	8.04E-5	4.97E-3	5.09E-4
900	8	2.40E-4	5.76E-3	2.67E-3
1000	6	3.22E-3	4.04E-3	4.54E-3
1100	4	4.82E-4	2.41E-4	5.03E-3
1200	2	9.63E-4	4.81E-4	1.16E-1

(a) The uncertainties for the mass flux measurements are given in the calculations in Appendix A.

4. DISCUSSION

We have plotted the mass flux of tantalum and tungsten with respect to temperature in Figure 13. The value for tungsten is approximately one-tenth of that for tantalum at 600°C whereas the mass flux of hafnium from Table 3 is about two orders of magnitude lower than tantalum. These ratios correlate with those in the alloy suggesting that mobilization from the specimen was by oxide spalling. There was evidence that tungsten was also mobilized by another mechanism involving volatilization. One-third of the tungsten mobilized at 500°C was found deposited downstream from the test chamber. About half of the W mobilized was found downstream of the test chamber at 800°C and about 85 percent at 1200°C. This higher relative migration of tungsten from the test chamber with increasing temperature and the visible evidence of a deposit on the filter from the 1200°C test show that migration was occurring by volatile species, i.e., WO_3 polymers, as previously noted. The decrease in tantalum mobilization with temperature correlate with the amounts of powder that we observed for the various tests. Most notably was the lack of spalling or oxide scale exfoliation at the highest temperatures of 1100 and 1200°C. These observations plus the fact that there was little indication of tantalum migration beyond the test chamber show that the particulate we characterized in Section 3.4 are not very susceptible to migration.

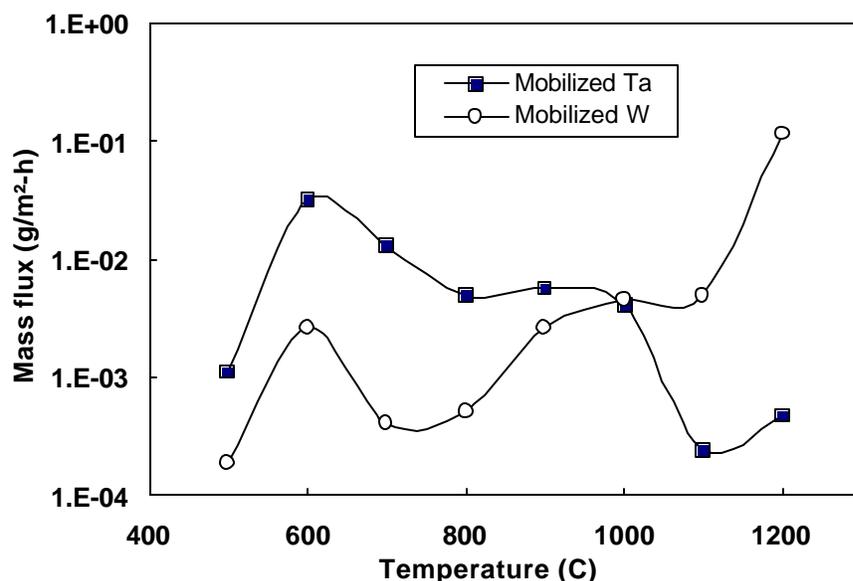


Figure 13. Mass flux of tantalum and tungsten from T-222 plotted with respect to temperature.

We compare the mass flux of tungsten and tantalum from the T-222 alloy with values from our previous study of air oxidation of a powder metallurgy (PM) product.[5] The PM product contained 95 and 0.1 wt% tungsten and tantalum, respectively. Figure 14 shows that the mass flux of tungsten for the two different studies was quite comparable at 500°C despite the fact that the T-222 alloy contained only one-tenth as much tungsten. The measurements from the T-222, however, are about two orders of magnitude lower above 800°C. This agrees with the

temperatures at which we observed the development and retention of significant oxide scale on T-222 alloy. Perhaps most of the WO_3 volatilization occurred during the initial stages of the test before an oxide scale formed. Once a scale formed similar to those shown in Figure 3, the volatilization process was impeded. Plots for tantalum mass flux in Figure 15 show a different trend. The values of tantalum mobilization for the PM product with only 0.1 wt% Ta exceed those from T-222 above 800°C. Considering the apparent link between tantalum mobilization and oxide spalling, results from the previous study[5] must be influenced by the spalling characteristics of the tungsten PM product.

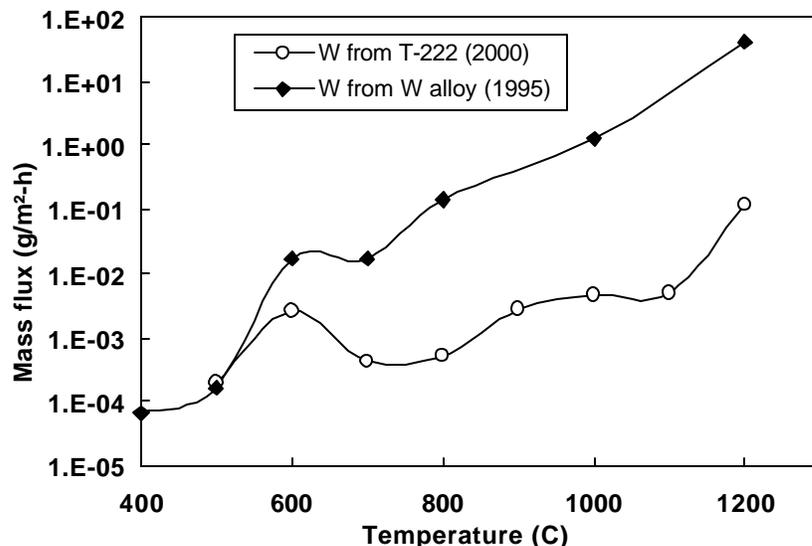


Figure 14. Mass flux of tungsten from T-222 alloy and a previously tested PM product containing 95 wt% tungsten.

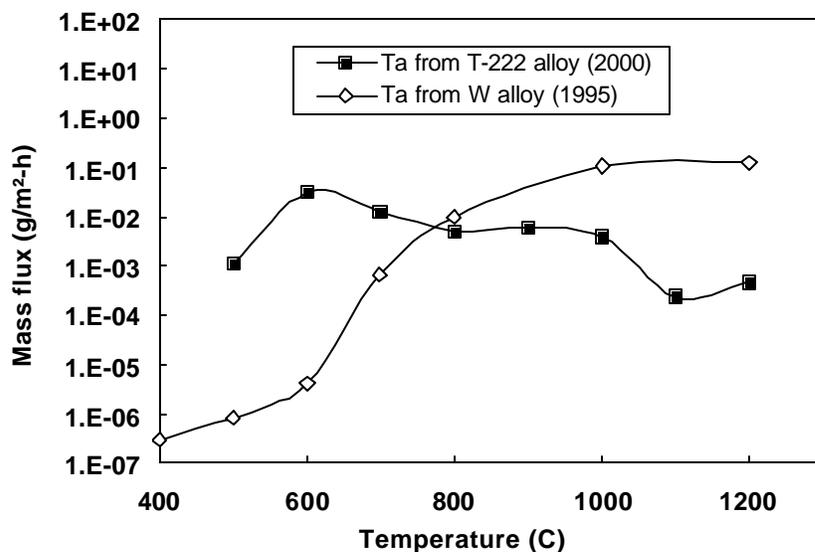


Figure 15. Mass flux of tantalum from T-222 alloy and a previously tested PM product containing only 0.1 wt% tantalum.

We observed embrittlement from oxygen dissolution into the tantalum alloy. These were apparent from the cracks and increased hardness in the alloy shown in Figure 10, Figure 11, and Figure 12. The specimen tested at 1100°C also fractured in a very brittle mode when we tried to remove the oxide scale. These observations show that there should be an awareness of potential embrittlement of tantalum alloys if exposed to significant oxygen pressures and temperatures during an off-normal condition.

5. CONCLUSIONS

We have found that mobilization processes in tantalum alloy T-222 is linked to the character of the oxide. The development and growth of oxides on tantalum are complex and depend upon time, temperature and oxygen pressure. Oxide spalling appears to be a major contributor to the mobilization process. We observed a smaller yellowish particulate that we believed formed during exposures. This type of spalled particulate and the measurements of mobilized tantalum and hafnium are most abundant at 600 and 700°C. Oxides appear to be sintered into a harder scale at higher temperatures. We believe that larger pieces of oxide scale and some particulate from near the oxide to metal interface are released during cool down. The oxide scales remained intact and very little particulate formation resulted from the highest temperature exposures at 1100 and 1200°C. Although the mobilization of tantalum and hafnium is basically linked to spalled particulate we did not observe the transport of even the small, flake-like particles out of the test chamber.

Tungsten was mobilized not only by spalling but showed an increasing contribution from volatilization with increasing temperature. Volatilization played a significant role even at the lowest temperature of 500°C. This may occur prior to the formation of the tantalum oxide scale. The mass flux at this temperature was actually very close to that which we observed from a previously tested powder metallurgy tungsten alloy. The volatilization of tungsten then actually decreases relative to its molar concentration in the alloy at higher temperatures. This suggests that an enhanced oxide scale slowed down the volatilization process of tungsten from the alloy.

We have determined mass flux values for tantalum, hafnium and tungsten from the T-222 alloy. These data could be used to estimate mobilization and activated product releases. However, since the processes are largely driven by oxidation spalling potential influences of thermal cycling with additional particulate formation should be kept in mind. We have also observed the effects of the dissolution of interstitial gas species into the tantalum alloy. This increased hardness and rendered some specimens quite brittle. This is an issue that should be considered in future applications of such alloys that could potentially be exposed to oxidizing conditions.

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APPENDIX A

**ICP-AES ANALYSES AND
MASS FLUX CALCULATIONS
FOR ELEMENTS MOBILIZED FROM
TANTALUM ALLOY T-222**

Ta1: Tested at 500°C for 81 hours in air (1000 sccm flow rate)

Specimen Ta1 (Date of Anal.)	Conditions	Component	Co wavelength	Hf wavelength	Ta wavelength	W wavelength
=====						
For Data analyzed on 9-19-00,		% SD of Cal. Stds	sigma %	3.2	5.4	4.2
9-27-00, 10-2-00, 10-09-		SD of Blk Std	0.002	0.020	0.007	0.006
Specimen						
Ta1	500C/81hours					
Air		RXN tube(set J)ug/ml	-0.012	-0.012	2.822	0.363
Time, h	05/15/2000	Col. tube,(Set 14) ug/ml	-0.043	-0.043	-0.011	0.058
		QW filter,ug/ml	0.011	0.011	-0.001	0.092
81	(A+B+C)	subtotal, ug	-2.200	-2.200	140.500	25.650
		± ug	0.147	0.147	15.239	1.592
	a (RXN)	Ta1 RXN FB, ug/ml	0.011	0.011	0.012	0.002
	b (Col T)	Ta1 CT FB, ug/ml	-0.001	-0.001	-0.013	0.023
	c (QW blank)	QW mean, ug/ml	0.030	0.030	0.001	0.012
	(a+b+c)	subtotal, ug	2.000	2.000	0.000	1.850
		± ug	3.464	3.464	1.212	1.039
	(A+B+C)-(a+b+c)	Total, ug	-4.200	-4.200	140.500	23.800
		± ug	3.467	3.467	15.287	1.901
	DL (3*SD)	ug/ml	0.060	0.060	0.021	0.018
	DL (3*SD)	ug	3.000	3.000	1.050	0.900
	Total Corrected for DL		3.000	3.000	140.500	23.800
						Total g/m2-h
	As calculated flux values:	Mass Flux, g/m2-h	2.38E-05	2.38E-05	1.12E-03	1.89E-04
		± g/m2-h	2.75E-05	2.75E-05	1.21E-04	1.51E-05
		± %	115.57	115.57	10.88	7.99
						1.67E-04
	Flux values with Eng.Judgement:	Mass Flux, g/m2-h	2.38E-05	2.38E-05	1.12E-03	1.89E-04
	Errors not applicable	± g/m2-h	2.75E-05	2.75E-05	1.21E-04	1.51E-05
		± %	115.57	115.57	10.88	7.99
=====						
Specimen: Ta1						
Dimensions:	t avg, cm	h avg., cm	w avg., cm			
	0.2690	2.5420	2.5232			
Area, m2	0.001555					
weight, g	28.6025					

Ta2: Tested at 600°C for 55 hours in air (1000 sccm flow rate)

Specimen Ta2 (Date of Anal.)	Conditions	Component	Co wavelength	Hf wavelength	Ta wavelength	W wavelength
			228.6	282.0	263.5	207.9
=====						
For Data analyzed on 9-19-00, 9-27-00, 10-2-00, 10-09-		sigma %	10.4	3.2	5.4	4.2
Specimen		sigma prime ug/ml	0.002	0.020	0.007	0.006
Ta2	600C/55 hours					
Air	A	RXN tube(no ID)ug/ml		0.482	53.410	4.544
Time, h	05/23/2000	Col. tube.(Set 12) ug/ml		-0.034	-0.008	0.016
	C	QW filter,ug/ml		-0.024	0.034	-0.002
55	(A+B+C)	subtotal, ug		21.200	2671.800	227.900
		± ug		1.548	288.414	19.085
	a (RXN)	Ta2 RXN FB, ug/ml		-0.016	0.007	0.016
Notes:	b (Col T)	Ta2 CT FB, ug/ml		-0.014	-0.005	0.013
	c (QW blank)	QW mean, ug/ml		0.030	0.001	0.012
1) All 50 ml aloquits.	(a+b+c)	subtotal, ug		0.000	0.150	2.050
		± ug		3.464	1.212	1.039
	(A+B+C)-(a+b+c)	Total, ug		21.200	2671.650	225.850
		± ug		3.794	288.417	19.113
	DL (3*SD)	ug/ml		0.060	0.021	0.018
	DL (3*SD)	ug		3.000	1.050	0.900
	Total Corrected for DL			21.200	2671.650	225.850
						Total g/m2-h
As calculated flux values:		Mass Flux, g/m2-h		2.48E-04	3.12E-02	2.64E-03
		± g/m2-h		4.44E-05	3.37E-03	2.23E-04
		± %		17.90	10.80	8.46
Flux values with Eng.Judgement:		Mass Flux, g/m2-h		2.48E-04	3.12E-02	2.64E-03
Errors not applicable		± g/m2-h		4.44E-05	3.37E-03	2.23E-04
		± %		17.90	10.80	8.46
Specimen: Ta2						
Dimensions:	t avg, cm	h avg., cm	w avg., cm			
	0.2694	2.5415	2.5225			
Area, m2	0.001555					
weight, g	28.8577					

Ta3: Tested at 700°C for 55 hours in air (1000 sccm flow rate)

Specimen Ta3 (Date of Anal.)	Conditions	Component	Co wavelength	Hf wavelength	Ta wavelength	W wavelength
			228.6	282.0	263.5	207.9
=====						
For Data analyzed on 9-19-00,						
9-27-00, 10-2-00, 10-09-		sigma %	10.4	3.2	5.4	4.2
Specimen		sigma prime ug/ml	0.002	0.020	0.007	0.006
Ta3	700C/55 hours					
Air		RXN tube(no ID)ug/ml		0.177	22.260	0.444
Time, h	05/30/2000	Col. tube,(Set 12) ug/ml		-0.010	0.003	0.268
		QW filter,ug/ml		0.029	0.025	0.025
55		subtotal, ug		9.800	1114.400	36.850
		± ug		0.575	120.204	2.181
		Ta3 RXN FB, ug/ml		0.059	0.021	0.009
Notes:		Ta3 CT FB, ug/ml		-0.015	-0.006	0.014
		QW mean, ug/ml		0.030	0.001	0.012
1) All 50 ml aloqois.		subtotal, ug		3.700	0.800	1.750
		± ug		3.464	1.212	1.039
		Total, ug		6.100	1113.600	35.100
		± ug		3.511	120.210	2.416
		ug/ml		0.060	0.021	0.018
		ug		3.000	1.050	0.900
		Total Corrected for DL		6.100	1113.600	35.100
						Total g/m2-h
		Mass Flux, g/m2-h		7.13E-05	1.30E-02	4.10E-04
As calculated flux values:		± g/m2-h		4.10E-05	1.40E-03	2.82E-05
		± %		57.57	10.79	6.88
		Mass Flux, g/m2-h		7.13E-05	1.30E-02	4.10E-04
Flux values with Eng.Judgement:		± g/m2-h		4.10E-05	1.40E-03	2.82E-05
Errors not applicable		± %		57.57	10.79	6.88
Specimen: Ta3						
		t avg, cm				
Dimensions:		h avg, cm	w avg, cm			
		0.2694	2.5433			
Area, m2		0.001556				
weight, g		28.7979				

Ta4: Tested at 800°C for 48 hours in air (1000 sccm flow rate)

Specimen Ta4 (Date of Anal.)	Conditions	Component	Co wavelength	Hf wavelength	Ta wavelength	W wavelength
=====						
For Data analyzed on 9-19-00,						
9-27-00, 10-2-00, 10-09-	% SD of Cal. Stds	sigma %	10.4	3.2	5.4	4.2
Specimen	SD of Blk Std	sigma prime ug/ml	0.002	0.020	0.007	0.006
Ta4	800C/48 hours					
Air	A	RXN tube(no ID)ug/ml		0.045	7.394	0.391
Time, h	08/01/2000	B	Col. tube,(Set 12) ug/ml	0.006	0.005	0.043
		C	QW filter,ug/ml	0.037	0.019	0.370
48	(A+B+C)	subtotal, ug	4.400	370.900	40.200	
		± ug	0.187	39.928	2.268	
		Ta4 RXN FB, ug/ml	-0.021	0.002	0.018	
Notes:	b (Col T)	Ta4 CT FB, ug/ml	-0.041	0.001	0.014	
	c (QW blank)	QW mean, ug/ml	0.030	0.001	0.012	
1) All 50 ml aloquits.	(a+b+c)	subtotal, ug	-1.600	0.200	2.200	
		± ug	3.464	1.212	1.039	
	(A+B+C)-(a+b+c)	Total, ug	6.000	370.700	38.000	
		± ug	3.469	39.946	2.495	
	DL (3*SD)	ug/ml	0.060	0.021	0.018	
	DL (3*SD)	ug	3.000	1.050	0.900	
	Total Corrected for DL		6.000	370.700	38.000	Total
						g/m2-h
As calculated flux values:		Mass Flux, g/m2-h		8.04E-05	4.97E-03	5.09E-04
		± g/m2-h		4.65E-05	5.35E-04	3.34E-05
		± %		57.82	10.78	6.57
						g
						4.15E-04
Flux values with Eng.Judgement:		Mass Flux, g/m2-h		8.04E-05	4.97E-03	5.09E-04
Errors not applicable		± g/m2-h		4.65E-05	5.35E-04	3.34E-05
		± %		57.82	10.78	6.57
Specimen: Ta4						
Dimensions:	t avg, cm	h avg., cm	w avg., cm			
	0.2694	2.5232	2.5408			
Area, m2	0.001555					
weight, g	28.8238					

Ta5: Tested at 900°C for 8 hours in air (1000 sccm flow rate)

Specimen Ta5 (Date of Anal.)	Conditions	Component	Co wavelength	Hf wavelength	Ta wavelength	W wavelength
=====						
For Data analyzed on 9-19-00,						
9-27-00, 10-2-00, 10-09-	% SD of Cal. Stds	sigma %	10.4	3.2	5.4	4.2
Specimen	SD of Blk Std	sigma prime ug/ml	0.002	0.020	0.007	0.006
Ta5	900C/8 hours					
Air	A	RXN tube(no ID)ug/ml	0.017	0.017	1.435	0.168
Time, h	08/08/2000	Col. tube,(Set 12) ug/ml	0.018	0.018	0.005	0.155
	C	QW filter,ug/ml	0.031	0.031	0.001	0.355
8	(A+B+C)	subtotal, ug	3.300	3.300	72.050	33.900
		± ug	0.127	0.127	7.749	1.773
	a (RXN)	Ta5 RXN FB, ug/ml	0.021	0.021	0.003	-0.003
Notes:	b (Col T)	Ta5 CT FB, ug/ml	-0.015	-0.015	-0.001	0.001
	c (QW blank)	QW mean, ug/ml	0.030	0.030	0.001	0.012
1) All 50 ml aloquits.	(a+b+c)	subtotal, ug	1.800	1.800	0.150	0.500
		± ug	3.464	3.464	1.212	1.039
	(A+B+C)-(a+b+c)	Total, ug	1.500	1.500	71.900	33.400
		± ug	3.466	3.466	7.843	2.055
	DL (3*SD)	ug/ml	0.060	0.060	0.021	0.018
	DL (3*SD)	ug	3.000	3.000	1.050	0.900
	Total Corrected for DL		3.000	3.000	71.900	33.400
						Total g/m2-h 8.67E-03
As calculated flux values:		Mass Flux, g/m2-h	2.40E-04	2.40E-04	5.76E-03	2.67E-03
		± g/m2-h	2.78E-04	2.78E-04	6.28E-04	1.65E-04
		± %	115.55	115.55	10.91	6.15
						g 1.08E-04
Flux values with Eng.Judgement:		Mass Flux, g/m2-h	2.40E-04	2.40E-04	5.76E-03	2.67E-03
Errors not applicable		± g/m2-h	2.78E-04	2.78E-04	6.28E-04	1.65E-04
		± %	115.55	115.55	10.91	6.15
=====						
Specimen: Ta5						
Dimensions:	t avg, cm	h avg., cm	w avg., cm			
	0.2696	2.5263	2.5476			
Area, m2	0.001561					
weight, g	28.9162					

Ta6: Tested at 1000°C for 6 hours in air (1000 sccm flow rate)

Specimen Ta6 (Date of Anal.)	Conditions	Component	Co wavelength	Hf wavelength	Ta wavelength	W wavelength
=====						
For Data analyzed on 9-19-00, 9-27-00, 10-2-00,10-09-		sigma %	10.4	3.2	5.4	4.2
SD of Blk Std		sigma prime ug/ml	0.002	0.020	0.007	0.006
Specimen						
Ta6	1000C/6 hours					
Air		RXN tube(no ID)ug/ml		0.010	0.723	0.536
Time, h	08/10/2000	Col. tube,(Set 12) ug/ml		-0.030	0.040	0.018
		QW filter,ug/ml		-0.001	-0.001	0.325
6	(A+B+C)	subtotal, ug		-1.050	38.100	43.950
		± ug		0.101	3.910	2.634
	a (RXN)	Ta6 RXN FB, ug/ml		0.045	0.001	0.006
Notes:	b (Col T)	Ta6 CT FB, ug/ml		0.005	0.007	0.014
	c (QW blank)	QW mean, ug/ml		0.030	0.001	0.012
	(a+b+c)	subtotal, ug		4.000	0.450	1.600
		± ug		3.464	1.212	1.039
	(A+B+C)-(a+b+c)	Total, ug		-5.050	37.650	42.350
		± ug		3.466	4.094	2.831
	DL (3*SD)	ug/ml		0.060	0.021	0.018
	DL (3*SD)	ug		3.000	1.050	0.900
	Total Corrected for DL			3.000	37.650	42.350
						Total g/m2-h
	As calculated flux values:	Mass Flux, g/m2-h		3.22E-04	4.04E-03	4.54E-03
		± g/m2-h		3.72E-04	4.39E-04	3.04E-04
		± %		115.52	10.87	6.69
						8.30E-05
	Flux values with Eng.Judgement:	Mass Flux, g/m2-h		3.22E-04	4.04E-03	4.54E-03
	Errors not applicable	± g/m2-h		3.72E-04	4.39E-04	3.04E-04
		± %		115.52	10.87	6.69
Specimen: Ta6						
Dimensions:	t avg, cm	h avg., cm	w avg., cm			
	0.2692	2.5235	2.5377			
Area, m2	0.001553					
weight, g	28.8064					

Ta7: Tested at 1100°C for 4 hours in air (1000 sccm flow rate)

Specimen Ta7 (Date of Anal.)	Conditions	Component	Co wavelength	Hf wavelength	Ta wavelength	W wavelength
=====						
For Data analyzed on 9-19-00,						
9-27-00, 10-2-00, 10-09-	% SD of Cal. Stds	sigma %	10.4	3.2	5.4	4.2
Specimen	SD of Blk Std	sigma prime ug/ml	0.002	0.020	0.007	0.006
Ta7	1100C/4 hours					
Air	A	RXN tube(no ID)ug/ml		-0.038	0.035	0.660
Time, h	08/15/2000	B Col. tube,(Set 12) ug/ml		-0.008	-0.002	0.007
	C	QW filter,ug/ml		-0.021	0.008	0.030
4	(A+B+C)	subtotal, ug		-3.350	2.050	34.850
		± ug		0.141	0.194	2.775
	a (RXN)	Ta7 RXN FB, ug/ml		0.018	0.010	0.037
Notes:	b (Col T)	Ta7 CT FB, ug/ml		0.011		0.022
	c (QW blank)	QW mean, ug/ml		0.030	0.001	0.012
1) All 50 ml aloquits.	(a+b+c)	subtotal, ug		2.950	0.550	3.550
		± ug		3.464	1.212	1.039
	(A+B+C)-(a+b+c)	Total, ug		-6.300	1.500	31.300
		± ug		3.467	1.228	2.963
	DL (3*SD)	ug/ml		0.060	0.021	0.018
	DL (3*SD)	ug		3.000	1.050	0.900
	Total Corrected for DL			3.000	1.500	31.300
						Total g/m2-h
As calculated flux values:		Mass Flux, g/m2-h		4.82E-04	2.41E-04	5.03E-03
		± g/m2-h		5.57E-04	1.97E-04	4.76E-04
		± %		115.57	81.86	9.47
Flux values with Eng.Judgement:		Mass Flux, g/m2-h		4.82E-04	2.41E-04	5.03E-03
Errors not applicable		± g/m2-h		5.57E-04	1.97E-04	4.76E-04
		± %		115.57	81.86	9.47
Specimen: Ta7						
Dimensions:	t avg, cm	h avg., cm	w avg., cm			
	0.2696	2.5410	2.5254			
Area, m2	0.001557					
weight, g	28.8799					

Ta8: Tested at 1200°C for 2 hours in air (1000 sccm flow rate)

Specimen Ta8 (Date of Anal.)	Conditions	Component	Co wavelength	Hf wavelength	Ta wavelength	W wavelength
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For Data analyzed on 9-19-00,						
9-27-00, 10-2-00, 10-09-	% SD of Cal. Stds	sigma %	10.4	3.2	5.4	4.2
Specimen	SD of Blk Std	sigma prime ug/ml	0.002	0.020	0.007	0.006
Ta8	1200C/2 hours					
Air	A	RXN tube(no ID)ug/ml		-0.026	0.019	1.186
Time, h	08/17/2000	B	Col. tube,(Set 12) ug/ml	0.012	-0.009	0.200
		C	QW filter,ug/ml	-0.039	0.028	5.926
2	(A+B+C)	subtotal, ug		-2.650	1.900	365.600
		± ug		0.155	0.189	25.397
		Ta8 RXN FB, ug/ml		-0.013	0.000	0.021
Notes:		Ta8 CT FB, ug/ml		-0.038	0.007	0.021
		QW mean, ug/ml		0.030	0.001	0.012
1) All 50 ml aloquits.	(a+b+c)	subtotal, ug		-1.050	0.400	2.700
		± ug		3.464	1.212	1.039
	(A+B+C)-(a+b+c)	Total, ug		-1.600	1.500	362.900
		± ug		3.468	1.227	25.418
	DL (3*SD)	ug/ml		0.060	0.021	0.018
	DL (3*SD)	ug		3.000	1.050	0.900
	Total Corrected for DL			3.000	1.500	362.900
						Total g/m2-h
As calculated flux values:		Mass Flux, g/m2-h		9.63E-04	4.81E-04	1.16E-01
		± g/m2-h		1.11E-03	3.94E-04	8.16E-03
		± %		115.59	81.81	7.00
						g 3.67E-04
Flux values with Eng.Judgement:		Mass Flux, g/m2-h		9.63E-04	4.81E-04	1.16E-01
Errors not applicable		± g/m2-h		1.11E-03	3.94E-04	8.16E-03
		± %		115.59	81.81	7.00
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Specimen: Ta8						
Dimensions:	t avg, cm	h avg., cm	w avg., cm			
	0.2701	2.5450	2.5226			
Area, m2	0.001558					
weight, g	28.9705					