

# HIGH TEMPERATURE OXIDATION OF TUNGSTEN AND MOLYBDENUM

N. M. Ghoniem

The University of California at Los Angeles (UCLA) Los Angeles, CA. 90095-1597, USA

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### PRESENTATION OUTLINE

- (1) IMPACT OF OXIDATION ON DESIGN WITH REFRACTORY ALLOYS.
- (2) MODELING OF HIGH-TEMPERATURE OXIDATION.
  - 2.1. Thermodynamics of Reactions
  - 2.2. Non-equilibrium Analysis
- (3) RESULTS FOR TUNGSTEN.
- (4) RESULTS FOR MOLYBDENUM.
- (5) CONCLUSIONS.



### **DESIGN IMPACT**

- **@** At Normal temperatures and pressures, the chemical reaction of a gas with the solid generally results in condensed products.
- @ At high temperatures and low pressures, the formation of volatile products is thermodynamically favored over the growth of the condensed phase.
- @ The upper temperature limit for design with refractory metals with a helium coolant will be influenced by the formation of volatile oxides.
  - @ The present investigation is concerned with W/He and Mo/He designs.



### **MODELING HIGH-TEMPERATURE OXIDATION**

### **APPROACH:**

- @ Quasi-equilibrium Treatment of Heterogeneous Reactions for the systems: O-W and O-Mo.
- @ Rate Limiting Step is the Trapping (adsorption) of Oxygen atoms until equilibrium.
- @ Assume that the helium pressure is  $P_{sys}$ , and its temperature  $T^*$ . The oxygen partial pressure is given by:

$$P_{O_2'} = appm \times 10^{-6} P_{sys}$$



#### THERMODYNAMICS OF CHEMICAL REACTIONS

**@** The rate at which oxygen molecules collide with unit area of the wall surface is given by:

$$Z_{O_2} = P_{O_2} \sqrt{2\pi} \, M_{O_2} RT$$

**@** For Tungsten, the following thermodynamic reactions take place:

$$x W(s) + \frac{1}{2} y O_2(g) \Leftrightarrow W_x O_y(g)$$

@ For Molybdenum, similar reactions occur. In addition, for  $T\sim1500-2500~\mathrm{K}$ :

$$Mo(s) + \frac{1}{2}(3-\alpha) O_2(g) \Leftrightarrow Mo O_2(g) + (1-\alpha)Mo O_3(g)$$



### FORMATION ENTHALPIES & ENTROPIES OF OXIDES

Tungsten		Molybdenum		
Species	DHf <sub>298.15</sub> (kcal/g.mole)	Species	DHf <sub>298.15</sub>	DSf <sub>298.15</sub>
_		_	(kcal/g.mole)	(cal/g.mole.K)
$\mathbf{O}(\mathbf{g})$	59.559	O(g)	61.3	16
$\mathbf{W} \mathbf{O}(\mathbf{g})$	101.6	Mo O(g)	95	25.5
$WO_2(g)$	18.3	$MoO_{2}\left( g\right)$	11.0	9.0
$W O_3(g)$	-70.0	<b>Mo O</b> <sub>3</sub> (g)	-80	-15.5
$W_2 O_6 (g)$	-278.2	$Mo_2 O_6 (g)$	-270	-72
$W_3O_8(g)$	-408.7	$Mo_3O_8(g)$	-400	-118
$W_3O_9(g)$	-483.6	$Mo_3O_9(g)$	-463	-132
$W_4O_{12}(g)$	-670.2	$Mo_4O_{12}(g)$	-640	-190



# QUASI-EQUILIBRIUM TREATMENT OF BATTY & STICKNEY\*

- @ Collisions of  $O_2$  with the surface can lead to either: (1) adsorption and equilibration, or (2) reflection.
- @ Define the equilibration probability as:

$$\zeta_i = \Gamma_i / Z_i$$

@ For oxygen molecules at a temperature  $T^*$ , different from the wall temperature T, the equilibrated oxygen flux is:

$$\Gamma_{O_2'} = \zeta_{O_2'} Z_{O_2'}$$

J.C. Batty and R.E. Stickney, Journ. Chem. Physics, Volume 51, No. 10, (1969) p 4475



# QUASI-EQUILIBRIUM TREATMENT OF BATTY & STICKNEY\* (Cont.)

- @ Let  $i = W_x O_y(g)$ , and the Gibbs Energy as:  $\Delta G_i(T) = \Delta H_i(T) T\Delta S_i(T)$
- @ Now, we have the following system:

$$K_{i} = \frac{P_{i}}{(P_{O_{2}})^{y/2}} = \exp(-\Delta G_{i}(T) / RT) \qquad i = 1, 2, ..., N$$

$$P_{O} = \sqrt{P_{O_{2}}} \exp(-\Delta G_{O} / RT)$$

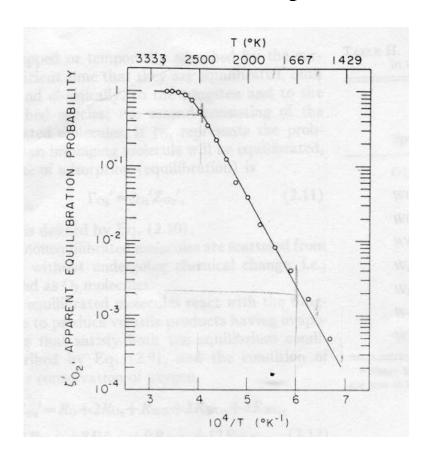
$$P_{O_{2}'} = P_{O_{2}} + P_{O} + \sum_{1}^{N} P_{i}$$

@  $P_{O2}$ ' is obtained from  $G_{O2}$ '. Requires equilibration probability

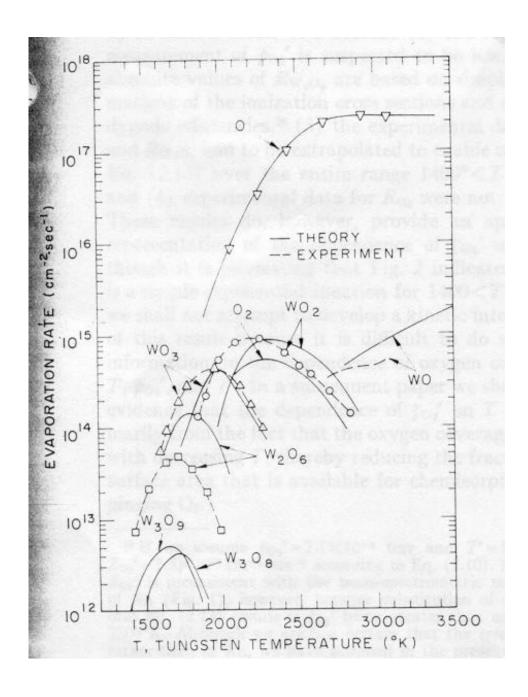


# EQUILIBRATION PROBABILITY IS DETERMINED EXPERIMENTALLY

$$\zeta_{o_2} = \exp[10.3498 - \frac{2.7607 \times 10^4}{T}]$$
 , T (°K)

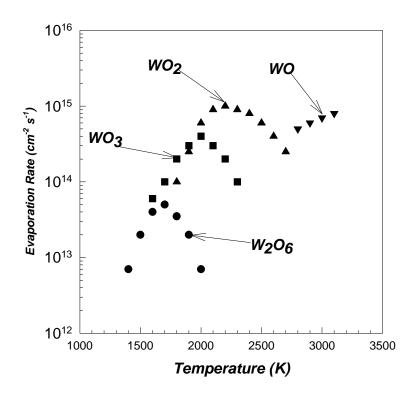






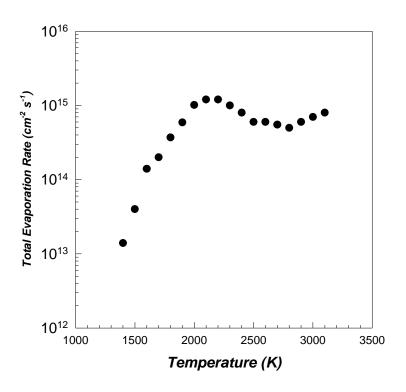


### Experimental Data on W Oxidation $Z_{02}$ '=1.2x10<sup>17</sup> cm<sup>-2</sup> s<sup>-1</sup>



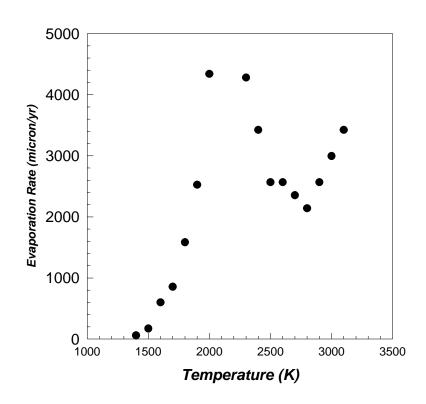


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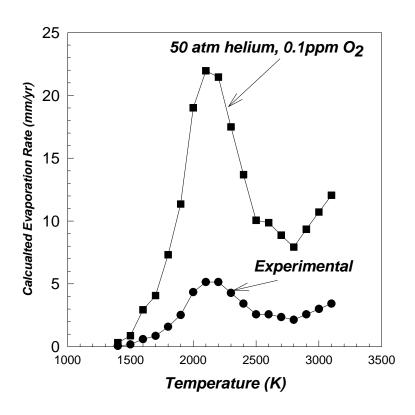


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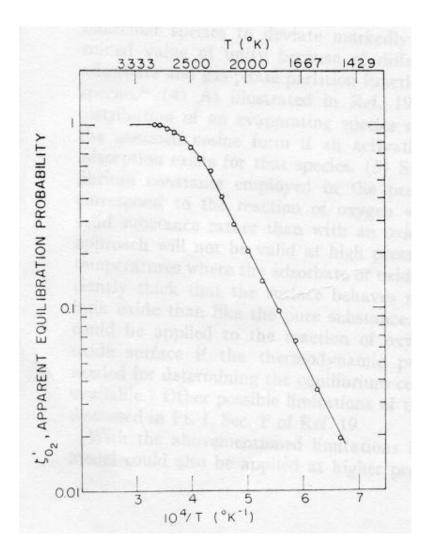


### Calculated W Oxidation Rates Experiment at $Z_{02}$ '=1.2x10<sup>17</sup> cm<sup>-2</sup> s<sup>-1</sup>

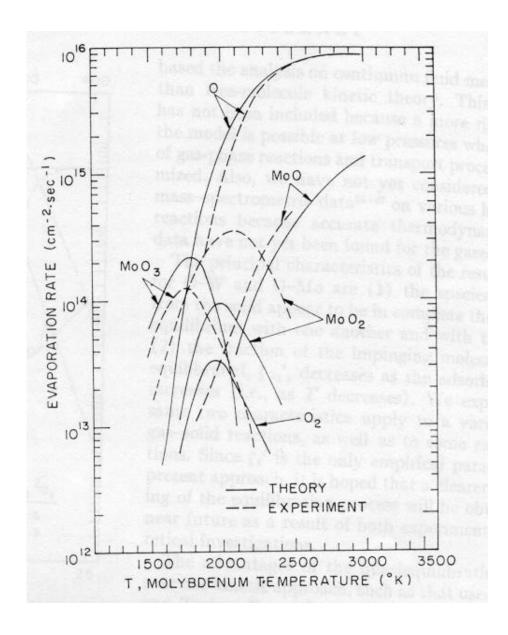




### EXPERIMENTAL RESULTS FOR MOLYBDENUM









### **CONCLUSIONS**

- (1) HIGH-TEMPERATURE OXIDATION SETS SEVERE LIMITS ON HELIUM COOLING OF REFRACTORY ALLOYS.
- (2) FOR TUNGSTEN AND MOLYBDENUM OPERATING AT 50 ATM. HELIUM COOLANT, AT 0.1 PPM OXYGEN, THE UPPER TEMPERATURE IS ESTIMATED AT 1200- 1300 °C.
- (3) IF HIGHER TEMPERATURES ARE BENIFICIAL TO DESIGN OBJECTIVES, THE FOLLOWING IS TO BE IMPLEMENTED:
  - @ Reduce the oxygen impurity concentration to the ppp range.
  - @ use oxidation-resistant coatings in the high-temperature zones.