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ADVANCES IN CERAMICS • VOLUME 27

FABRICATION AND PROPERTIES OF LITHIUM CERAMICS II

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HTO(g), or T₂O(g) from oxidic ceramic tritium breeders. The degree of surface coverage at a given time, the result of adsorption of H₂O or H₂, and diffusion of T₂ to the surface will determine which type of site is dominant in the release process and what the desorption activation energy will be at that time. Theoretical predictions of release curves by a model based on this desorption behavior have been successful in matching observed data.

References

observed release under steady state and transient conditions, a preliminary benchmark analysis has been carried out for the metasilicate sample P1 irradiated in the LISA-1 experiment. Initial results of this analysis are presented and the effect of several physical properties which potentially control the release kinetics are hereafter discussed.

Modeling Development

The Model Geometry

The model is based on the assumption of interconnected porosity in which the pores are represented as straight cylindrical capillaries having an effective radius $r_p$ and length $L_{e}$. The unit cell in the model has been identified as the interconnected pore system and the cylindrical portion of the solid that surrounds it. In the present work the following expression for $L_e$ is used:

$$ (L_{e})^2 = \tau $$

where $L$ is the characteristic length and $\tau$ is the tortuosity factor, which, according to the model of Wakao and Smith, is given by:

$$ \tau = \frac{1}{\varepsilon_p} $$

where $\varepsilon_p$ is the porosity of the material.

The pore radius $r_p$ is given by the following expression:

$$ r_p = \frac{2}{(S_{BET} \rho \cdot (1-\varepsilon_p))} $$

where $\rho$ = solid breeder density (TD) (Kg-m$^{-3}$); $S_{BET}$ = specific surface area (m$^2$-Kg$^{-1}$).

Tritium-Transport Mechanisms Included in the Model

The proposed model provides for a multi-region calculation and includes several transport mechanisms, which are considered to be relevant to controlling physical processes for ceramic breeder materials. A schematic of the multiregion model is shown in Fig. 1. The tritium, after being generated from breeding reactions in the grain, diffuses (most likely in anionic form) to the grain edges and to the grain-boundary paths and reaches the bulk/pore interface where it desorbs through the pores, most likely as molecules, in combination with any oxygen or hydrogen present at the surface. Tritium and tritium-bearing species then diffuse through the pores to the purge flow, where they are convected outside the solid breeder region. Therefore, surface phenomena, such as adsorption from the pore and from the bulk, desorption to the pore and eventually dissolution into the bulk, can play an important role in the overall tritium release kinetics and strongly influence the amount of gas which resides on the surfaces (expressed in terms of fractional coverage). Because of the complex nature of the mechanisms involved and the lack of basic experimental data, the characterization of the surface adsorption/desorption processes on lithium-based ceramics is very difficult. The modeling takes explicit account of the surface and its coupling with the interconnected pore and the bulk. Figure 2 shows a schematic diagram of the basic surface processes included in the model. The rate theory has been used to model the surface processes and the details of the scheme adopted are hereafter outlined.

Formation of the Governing Equations

Grain and Grain Boundary Regions

The time-dependent diffusion equations for the tritium concentration within the grain and the grain boundary assuming spherical grains and slab-shaped grain boundaries can be written as follows:

Fig. 1. Schematic of the multi-region model.
The applicable boundary, interface and initial conditions in conjunction with Eqs. (1) and (2) are

\[ C_y(r = 0, t) = \text{finite} \]  
\[ C_y(r = a, t) = C_y(x = 0, t) \]  
\[ 4\pi a^2 \cdot D_y(T) \left[ \frac{\partial C_y}{\partial r} \right]_{r=a} = N_{gb} \cdot A_{gb} \cdot D_y(T) \left[ \frac{\partial C_y}{\partial x} \right]_{x=0} \]  
\[ A_{gb} \cdot D_y(T) \left[ \frac{\partial C_y}{\partial x} \right]_{x=b} = R_{bulk} C_b(x = b, t)(1 - \theta) \]  
\[ C_y(r, t = 0) = C_{gb}(r) \quad \text{for} \quad 0 \leq r \leq a \]  
\[ C_y(x, t = 0) = C_{bo}(x) \quad \text{for} \quad 0 \leq x \leq b \]  

In Eq. (5), \( A_{gb} \) is the transversal area of the grain boundary path and \( N_{gb} \) is the effective number of one-dimension grain boundary paths associated with each grain. In Eq. (6), which couples the bulk region with the surface region, the term on the right represents the adsorption flux from the bulk. \( R_{bulk} \) is the rate constant for adsorption from the bulk, \( C_y(x = b, t) \) is the concentration in the grain boundary, just beneath the surface, and \( (1 - \theta) \) represents the fraction of sites available for atoms coming from the bulk to be adsorbed on the surface. The right side of terms of Eqs. (7) and (8) are the initial tritium concentrations within the grain and grain boundary, respectively.

**Surface Region**

The three fluxes going to and from the surface shown in Fig. 2 are included in our surface model.\(^4\) Balancing the adsorption fluxes from the pore and from the bulk with the desorption flux to the pore yields the following rate equation governing the coverage.
\[
\frac{d[N_j \cdot \theta_j]}{dt} = R_{\text{ads}}(T) \cdot C_{p}^0(z,t)(1-\theta)^2 + R_{\text{bulk}}(T)C_{b}(x=b,t)(1-\theta) - R_{\text{des}}(T) \cdot \theta_j \theta
\]

where \(\theta_j\) is fractional coverage for the adsorbed species \(j\) (i.e., atomic fraction of tritium or hydrogen atoms on the surface); \(\theta = total\ coverage = \Sigma \theta_j\); \(N_s = number\ of\ sites\ available\ (-10^{10}\ sites/m^2);\ C_{p}^0(z,t) = concentration\ of\ species\ (i)\ along\ the\ pore\ at\ spatial\ coordinate\ z,\ and\ time\ t\ (molecules\ (i)-m^{-2})\ ((i)\ could\ represent\ either\ H_2,\ T_2,\ HT,\ or\ H_2O,\ T_2O,\ and\ HTO);\ and\ the\ rate\ constant\ can\ be\ expressed\ as\ follows:\)

\[
R_{\text{ads}}(T) = \frac{\alpha}{\sqrt{8 \times 10^{-3} M_{(i)}}} \cdot \sqrt{RT} \cdot \exp(-E_{\text{ads}/RT})
\]

\[
R_{\text{des}}(T) = \nu_0 N_s \exp(-E_{\text{des}/RT})
\]

\[
R_{\text{bulk}}(T) = \beta \exp(-E_{\mu/RT})
\]

where \(E_{\text{ads}}, E_{\text{des}}, E_{\mu}\) are activation energies, respectively, for adsorption from the pore and from the bulk and for desorption from the surface, and \(\nu_0\) and \(\beta\) are pre-exponential of the order of \(1 \times 10^{13} \text{ s}^{-1}\).

**Interconnected Pore Region**

The pore is modeled as a straight cylinder and the governing 1-D diffusion equation can be written as follows for each tritium-bearing species.

\[
\frac{\partial C_{p}^0(z,t)}{\partial t} = D_{\text{eff}}^0(T) \frac{\partial^2 C_{p}^0(z,t)}{\partial z^2} + S_{p}^0(z,t)
\]

where \(D_{\text{eff}}^0(T)\) = effective diffusion coefficient along the pore for species (i) \((m^2/s)\); \(S_{p}^0(z,t)\) = volumetric source rate within the pore \((molecules\ (i)-m^{-3}s^{-1})\).

The boundary conditions for Eq. (13) are

\[
\frac{\partial C_{p}^0}{\partial z} \bigg|_{z=0} = 0
\]

(i.e., zero gradient at the beginning of the interconnected porosity system)

\[
C_{p}^0(z=L,t) = C_{p_0}
\]

where in Eq. (15) \(C_{p_0}\) represents the concentration of the species (i) gas in the purge stream, which is assumed known.

The source term in the pore diffusion equation couples the surface and the bulk with the interconnected pore system, and has to be expressed as the net rate of molecules leaving the surface at the solid/gas interface (i.e. desorption minus adsorption).

\[
S_{p}^0(z,t) = (R_{\text{des}}(T) \rho \cdot v_{\text{eff}} \cdot N_{\text{surf}}) \cdot \frac{S_{\text{surf}} \cdot \rho \cdot \nu}{V_{p}}
\]

where \(V_{p}\) is the volume of the interconnected porosity system and \(N_{\text{surf}}\) is the number of grains associated with one interconnected porosity system.

At the present stage, the analysis was performed for LISA-1 cases with only tritium in the system, so that in the previous set of equations

\[
\theta_j = \theta = \theta_T
\]

\[
C_{p}^0 = C_{p_2}
\]

The set of Eqs. (1) through (16) are solved numerically at each time step for each morphological breeder region (grain interior, grain boundary, surface, interconnected pore).

**Interconnected Pore Region**

\[
D_{\text{eff}}^0 = \varepsilon p^2 D_{p}^0
\]
\[ D_p^{(0)} = \frac{1}{1 - \alpha \cdot Y_i} + \frac{1}{D_o^{(0)}} \]

(20)

and the Knudsen diffusivity \( D_k^{(0)} \), which predominates when the characteristic length is of the order of the molecular mean free path, is given as:

\[ D_k^{(0)} \text{ (m}^2 \cdot \text{s}^{-1}) = 0.97 \tau_p \sqrt{\frac{T}{M_{W0}}} \]

(21)

for ordinary diffusion:

\[ D_e^{(0)} \text{ (m}^2 \cdot \text{s}^{-1}) = \frac{1.86 \times 10^{-7} \ T^{3/2}}{P \sigma^2 \Omega_i} \]

(22)

\[ M_{W0} = \text{molecular weight for species i} \]

\[ \alpha = 1 - \frac{M_{W0}}{M_{He}} \]

\[ Y_i = \text{fraction of species i; } P = \text{pressure in pore; } \sigma, \Omega_i = \text{constants described in Ref. 6.} \]

**Results of the Calculations and Discussion**

To test the proposed model quantitatively against available experimental data, we have chosen as a part of an agreed benchmarking program to examine some of the observed results for the LISA-1 experiment. For this paper, a detailed analysis of the release from the sample P1 (lithium metasilicate) under temperature transients, in the region of 600°-685°C with pure helium as purge gas is reported. These calculations were made with the data given in Table I and II, and the results are shown in the next set of figures. Figure 3 shows the calculated normalized tritium release profile from the end of the interconnected porosity system for the metasilicate sample P1. The temperature transient and the observed release profiles are also indicated. The model reproduces with
reasonably good accuracy the observed release profile and predicts quite well both the magnitude of the peaks and dips, produced by the corresponding temperature transients, and the slope of the profile which represents the rate of change of the release.

Previous modeling studies on the behavior of tritium transport during temperature transient conditions resulted in the identification of the ceramic breeder microstructure (e.g., grain size, pore size), the desorption activation energy, and the transient scenario as important factors affecting the release response. In this study the effects of the following key variables on the kinetics of the release, as well as on its relative magnitude, were investigated: (1) effect of the specific surface area, (2) effect of the desorption activation energy from the breeder surface, (3) effect of the diffusion activation energy within the grain, and (4) effect of gas diffusivity along the interconnected porosity system. Because of computing time constraints, the proposed parametric analyses were carried out over about half the total time interval shown in Fig. 3, precisely between 69.4 and 152.8 hrs, where effects due to both increase and decrease in temperature are contained.
In order to show the effect of the solid breeder specific surface area ($S_{BET}$) on tritium release response, a parametric analysis was carried out for three different values of $S_{BET}$: (1) 300, (2) 500, (3) 770 m$^2$/kg. The last value is given as reference for the orthosilicate. The resulting effect on tritium release is shown in Fig. 4. The best with the experimental profile is obtained for a value of $S_{BET} = 770$ m$^2$/kg, which was also used in Fig. 3. The profiles in Fig. 5 show the influence of the specific surface area on the tritium coverage. These results confirm the importance of the surface region, which acts like a valve and controls the rate of both filling of gas from the bulk and from the pore, as well as the desorption flux to the pore. Furthermore, to show how the magnitude of the activation energy for the desorption process may affect the overall release kinetics a parametric analysis was also performed for three different values of the heat of adsorption. In our model, estimates of the activation energy for desorption have been made from values of heat of adsorption as a function of the coverage available for the H$_2$O(g)-LiAlO$_2$ system, on the basis that the activation energy for desorption is equal to the sum of the heat of adsorption and the activation energy for adsorption. The heat of adsorption data are used here as an approximation because of the lack of data for the desorption activation energy as a function of coverage for lithium ceramics and hydrogen or water, whose experimental determination is again recommended here. In the calculations a constant value of 50 KJ/mol for the adsorption activation energy from the pore and 20 KJ/mol for adsorption from the bulk were used following the assumption that the adsorption activation energies are low. Changes of 10–20% in these values were found to weakly affect the results. The value resulting from the following formula for the desorption activation energy was used.

$$E_{des}(\theta) = E_{ad} + f \times Q(\theta) \quad (23)$$

where $Q(\theta)$ is the heat of adsorption as a function of the coverage shown in Table II and $f$ is a factor, which was set respectively, for the three cases investigated, equal to 0.5, 1, and 1.2. Results of this parametric analysis for both tritium release and coverage are shown in Figs. 6 and 7. As expected, the coverage is substantially affected by changes in the desorption activation energy. The tritium release transients are markedly lower for the case with lowest desorption activation energy as a result of the coverage being ten times lower than for the reference case. However the tritium release behavior is not as influenced for the case with the highest activation energy since the increase of coverage, as compared to the reference case, is only about a factor of 1.5. Figures 8 and 9 show the effect of a change in the interconnected pore diffusion.
coefficient on the tritium release from the end of the interconnected pore system and the surface coverage. Three cases were investigated: (1) the value of $D_{\text{p,eff}}$ given by Eq. (19); (2) $0.1 \times D_{\text{p,eff}}$; (3) $10 \times D_{\text{p,eff}}$. The pore diffusion coefficient strongly affects the release response and the effect on the tritium release peak is quite marked. For fast diffusion in the pore ($10 \times D_{\text{p,eff}}$), the tritium concentration in the pore is low and the resulting coverage is also low. Thus, temperature transients for that case have a lower effect on the tritium release. For slow diffusion in the pore ($0.1 \times D_{\text{p,eff}}$), the reverse occurs and the tritium release is markedly affected by temperature transients. This could represent a case where sintering would have significantly closed the pores and effectively slowed down the pore diffusion. Finally, an analysis was performed by changing the value of the activation energy for diffusion within the grain. No relevant changes in the tritium release response were observed, showing that, as inferred by the above analysis, the tritium release behavior is not diffusion controlled, but surface controlled for the case with no hydrogen in the helium purge.

**Summary**

A comprehensive model for predicting transient tritium behavior in lithium-based ceramic materials has been developed. It represents an attempt to develop a computer code with an efficient predictive capability to describe the tritium transport and release kinetics as a function of different variables and for a wide range of operating conditions. The model follows the basic physics of the expected transport process sequence, and particular emphasis is given to the analysis of surface phenomena, such as gas adsorption and desorption, which seem to play a crucial role for the accurate prediction of the release kinetics and gas concentration distributions. It is important for a code to properly describe the underlying physics, since fits obtained with physically unsound models to experimental data by varying suitable effective parameters, will fail whenever extrapolations are needed to new ranges of operating conditions. The model was developed so as to include the capability of analyzing temperature and purge flow chemistry transients. In this paper, the following key issues are addressed in conjunction with the analysis of the metasilicate sample PI irradiated in the LISA-1 experiment: (1) identification of the transport steps relevant to reproduce the experimental tritium release profile and (2) identification of the key parameters which affect the release kinetics. The most important results from the application of the model for a temperature change under the condition of pure helium purge flow are the sensitivity of the tritium release behavior to crucial factors such as specific surface area, desorption activation energy, and pore diffusivity, which all influence the dynamics of tritium adsorption and desorption and determine the amount of tritium which resides on the surfaces.
This strongly indicates that tritium release is surface controlled in this case. It then becomes important to experimentally determine the values of the activation energies for adsorption and desorption as a function of the coverage for both the lithium ceramic/hydrogen and lithium ceramic/water cases. Future applications of the model will include analysis of experimental results for temperature transients with hydrogen in the purge and purge flow chemistry transients.

Acknowledgement

This work was performed under U.S. Department of Energy Contract DE-FG03-86ER52123.

References


Hydrogen-Tritium Exchange on the Surface of Ceramic Breeders

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Exchange reactions of hydrogen isotopes on the surface of Li2O and LiAl2O3 were experimentally studied. At first, the breeder was loaded with tritiated water vapor. Then, adsorbed HTO was desorbed with He + H2 (10-10000 ppm) sweep gas. Chemical forms of desorbed tritium were HT and HTO, and the fraction of HT found to become larger for higher H2 concentration. Reaction rate for HT release found to be proportional to the amount of residual tritium and to the square root hydrogen concentration in the sweep gas. This infers that dissociation of hydrogen occurs on the surface.

Introduction

One of the major issues in developing a fusion reactor system is that tritium production must be self-sustaining. Tritium must be generated in sufficient quantities in high TBR and must be recovered in a short time. Short tritium residence time and this corresponds to low tritium inventories in the blanket. In order to estimate the tritium inventory with good accuracy, the tritium migration processes must be fully understood. However, until now, surface reaction—desorption and/or exchange reaction—on the solid breeder material has not been extensively studied, although it may sometimes cause major fraction for tritium inventory in the blanket. In this connection, we have been studying surface reactions by several methods. By in situ tritium release experiments (TTTEs) at the University of Tokyo, the released chemical form tritium and the residence time were found to be strongly influenced by the chemical composition of the sweep gas. When hydrogen gas was added to the helium sweep gas, the fraction of HT in the sweep gas increased with 1