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HELIUM BLISTERING OF CERAMIC COATINGS ON HASTELLOY X AND Nb-1% Zr[†]

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Abstract

The surface damage of insulating ceramic coatings on Hastelloy X and Nb-1% Zr held at room temperature and at 300°C has been studied for both 100 KeV and 250 KeV helium ion irradiation for a dose range from 3.7×10^{18} to 1×10^{19} ions cm^{-2} . Blisters were observed after room temperature irradiation with both 100 KeV and 250 KeV helium ions. However, for irradiation at 300°C no blisters could be observed. The sharp rise in the helium permeation with temperature, observed by others for some glasses and ceramics, is thought to be responsible for this behaviour. These results suggest that for the energy range studied helium blistering has a negligible surface erosion effect on such coatings if they are operated at temperatures above 300°C.

INTRODUCTION

Certain fusion reactor concepts require that the entire first wall or parts of it consist of electrically insulating material. For example, the first wall of the reference theta-pinch reactor (RTPR)¹ is a structure with an insulator on a metal backing. This design is necessary to electrically insulate the first wall metal against the emf which develops the implosion heating. The insulating layer serves to prevent electrical breakdown between the plasma and blanket segments during the short implosion heating stage. It is necessary that the dielectric and mechanical properties of this composite structure of the first wall be maintained during the pulsed mode operation of such a reactor for a reasonable lifetime. In the RTPR-design alumina has been specified as the electrically insulating liner of the first wall, and Nb-1% Zr as the first wall structural metal. In order to minimize the radiation damage in the insulating liner the design incorporated a neutral gas blanket near the wall.

For such theta-pinch type of fusion reactors different types of ceramic insulator coatings are being developed by Atomic International. It was the aim of these studies to determine the behaviour of such coatings under energetic helium bombardment at different target temperatures.

EXPERIMENTAL TECHNIQUES

Two different ceramic coatings on two different types of substrates, Nb-1% Zr and Hastelloy X, were supplied by D. W. Keefer, Atomic International. The coating on Hastelloy X (coating #13) consisted of a mixture of 54.3% SiO₂, 38.7% BaO, 7.0% Al₂O₃, and the coating on Nb-1% Zr (coating #14) was a mixture of 52.3% SiO₂, 40.5% BaO, 7.2% Al₂O₃. (The concentrations are given in wt%.) Details of preparation of these coatings can be found elsewhere.² The coatings on Hastelloy X substrates had thicknesses ranging from 71 to 130 μm and the

thickness of coatings on Nb-1% Zr substrates ranged from 62 to 99 μm . The thicknesses of these coatings are about 40 to 85 times larger than the projected range (R_p) of 250 keV $^4\text{He}^+$ in Silicon ($R_p \approx 1.53 \mu\text{m}$). The coatings were irradiated with 100 keV or 250 keV $^4\text{He}^+$ ions from a 2-MeV Van de Graaff accelerator in high vacuum at a total pressure of $\approx 5 \times 10^{-8}$ Torr. The flux of the incident helium ions was held at 1×10^{14} ions $\text{cm}^{-2} \text{sec}^{-1}$. For the determination of the actual dose for the irradiation of these surfaces comparison of the secondary electron emission for these surfaces vs. those for metal target surfaces were made. The total dose values given in this paper are correct within only $\approx 20\%$. Other details of the irradiations are similar to those of the irradiations of metal targets described earlier.³ No optical or chemical polishing of the targets was considered necessary but all the targets were cleaned in ultrasonic baths of trichloroethylene, acetone, distilled water and methanol, prior to irradiation. For irradiation at 300°C the targets were heated directly by ohmic heating and the target temperatures were measured by an infrared pyrometer. The target surfaces were examined prior to and after the irradiation with the aid of a Cambridge stereoscan S4-10 scanning electron microscope. The ceramic coatings were kept at 300°C during irradiation and during observation in the scanning electron microscope.

RESULTS AND DISCUSSION

Figure 1 shows scanning electron micrographs of blisters formed on coatings 13 (Figures 1a and 1b) and 14 (Figures 1c and 1d) during room temperature irradiation with 100 keV and 250 keV helium ions for a total dose of 3.7×10^{17} ions cm^{-2} . The appearance of dome shaped blisters in both ceramic coatings suggests that the coatings can be plastically deformed readily. This is quite different from the pits observed in other ceramic materials such as SiC after

helium ion irradiation.³ The ceramic coatings used in the present studies have an amorphous structure with very little crystallinity. For both coatings the majority of the blisters have diameters ranging from 0.6 to 3 μm for 100-keV He^+ ion irradiation, and ranging from 3 to 5 μm for 250-keV He^+ ion irradiation. In several regions clusters having about 2-6 interconnected blisters have formed. The blister density (i.e., the number of blisters per unit area) for both coatings is about the same, that is, the blister density is $(1.9 \pm 0.3) \times 10^6$ blisters cm^{-2} for both helium ion energies used and for the total dose of 3.7×10^{18} ions cm^{-2} . These results differ from those given in the summary⁴ of this paper, since a correction to the total dose value due to secondary electron emission had not been applied to it. Since the compositions of these two coatings are not greatly different, it is not surprising that the blister sizes are about the same. At a dose of 3.7×10^{18} ions cm^{-2} the blisters have not exfoliated to a measurable degree.

Figure 2 shows blisters formed at higher doses. The micrograph in figure 2 shows blisters formed on a coating of SiO_2 on Ni-10Cr for room temperature irradiation with 100-keV He^+ ions for a total dose of 1.1×10^{19} ions cm^{-2} . In this case the blisters appear to be more spherical and have increased in diameter (as compared to the lower dose micrograph). The blister diameters are in the range of 0.6 to 1.5 μm and the blister density is $(4.1 \pm 0.4) \times 10^6$ blisters cm^{-2} , a value which is about a factor of 3 higher than the one for the lower dose. The blisters are still interconnected in some regions.

In the case of the SiO_2 coating, figure 3 shows blisters on coating irradiated at room temperature with 250-keV He^+ ions for a total dose of 1×10^{19} ions cm^{-2} . The blisters appear to be more dome shaped and have

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1. T. G. Parker, Jr., editor, "Development of Glass Electrical Insulator for High Voltage Power Lines," *IEEE Trans. on Power Apparatus and Systems*, PAS-86, 1967, pp. 100-107.
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3. M. Kurihara and S. K. Das, "Erosion of Silicon Carbide Surfaces Under High Voltage Discharge," *IEEE Trans. on Power Apparatus and Systems*, PAS-86, 1967, pp. 100-107.
4. M. Kurihara and S. K. Das, "Erosion of Silicon Carbide Surfaces Under High Voltage Discharge," *IEEE Trans. on Power Apparatus and Systems*, PAS-86, 1967, pp. 100-107.
5. M. Kurihara and S. K. Das, "Erosion of Silicon Carbide Surfaces Under High Voltage Discharge," *IEEE Trans. on Power Apparatus and Systems*, PAS-86, 1967, pp. 100-107.
6. M. Kurihara and S. K. Das, "Erosion of Silicon Carbide Surfaces Under High Voltage Discharge," *IEEE Trans. on Power Apparatus and Systems*, PAS-86, 1967, pp. 100-107.
7. M. Kurihara and S. K. Das, "Erosion of Silicon Carbide Surfaces Under High Voltage Discharge," *IEEE Trans. on Power Apparatus and Systems*, PAS-86, 1967, pp. 100-107.
8. V. O. A. Tomase, *J. Appl. Phys.*, "Helium Diffusion through Glass" 32, 1309

FIGURE CAPTIONS

Fig. 1. Scanning electron micrographs (SEI's) of ceramic coatings after irradiation at room temperature with ${}^4\text{He}^+$ ions to a total dose of 1.7×10^{19} ions/cm²: (a) coating #13 (see text for composition) on Ti-6Al-4V irradiated with ${}^4\text{He}^+$, (b) coating #13 irradiated with ${}^4\text{He}^+$, (c) coating #13 (see text for composition) on Ti-6Al-4V irradiated with ${}^4\text{He}^+$, (d) coating #14 irradiated with 250-keV ${}^4\text{He}^+$.

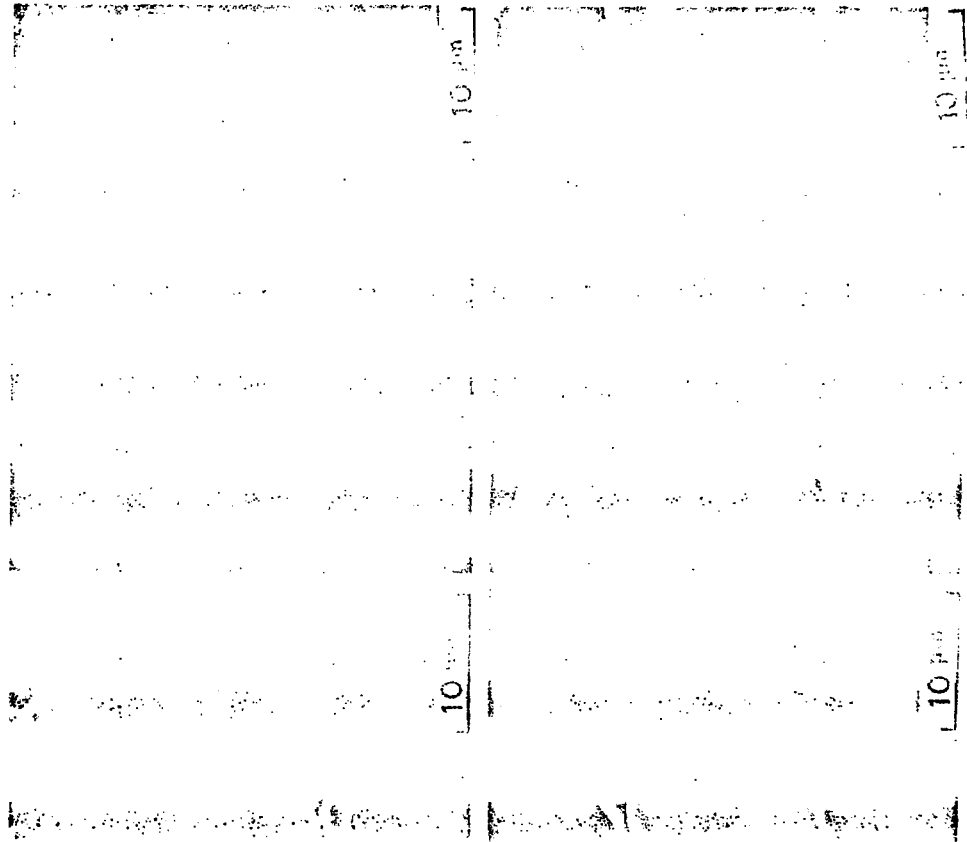
Fig. 2. SEMs of coating #14 on Nb-1% Zr irradiated at room temperature (a) with 100-keV ${}^4\text{He}^+$ ions to a dose of 1×10^{19} ions/cm², (b) with 250-keV ${}^4\text{He}^+$ ions to a dose of 1×10^{19} ions/cm², (c) with 250-keV ${}^4\text{He}^+$ ions to a total dose of 1×10^{19} ions/cm².

Fig. 3. SEMs of coating #14 on Nb-1% Zr irradiated at 600°C with 100-keV ${}^4\text{He}^+$ ions to a total dose of 3.7×10^{18} ions/cm², (a) coating #14 on Nb-1% Zr irradiated at 600°C with 100-keV ${}^4\text{He}^+$ ions to a total dose of 3.7×10^{18} ions/cm², (b) coating #14 on Hastelloy X irradiated at 600°C with 100-keV ${}^4\text{He}^+$ ions to a total dose of 3.7×10^{18} ions/cm², (c) coating #14 on Hastelloy X irradiated at 600°C with 100-keV ${}^4\text{He}^+$ ions to a total dose of 3.7×10^{18} ions/cm².

INSULATOR CERAMIC COATINGS IRRADIATED AT ROOM TEMP.
TO A DOSE OF 0.5 C/cm²

COATING no. 13 (54.3% SiO₂,
38.7% B₂O₃, 7.0% Al₂O₃) ON
HASTELLOY X

COATING no. 14 (54.3% SiO₂,
40.5% B₂O₃, 7.2% Al₂O₃) ON
ON Nb-1% Zr



WITH 100keV He⁺

WITH 250keV He⁺

CERAMIC COATING no.14 (52.5% SiO₂, 40.5% BaO, 7.2% Al₂O₃)
ON Nb-1%Zr IRRADIATED AT ROOM TEMP.

WITH 100keV He⁺

WITH 250keV He⁺

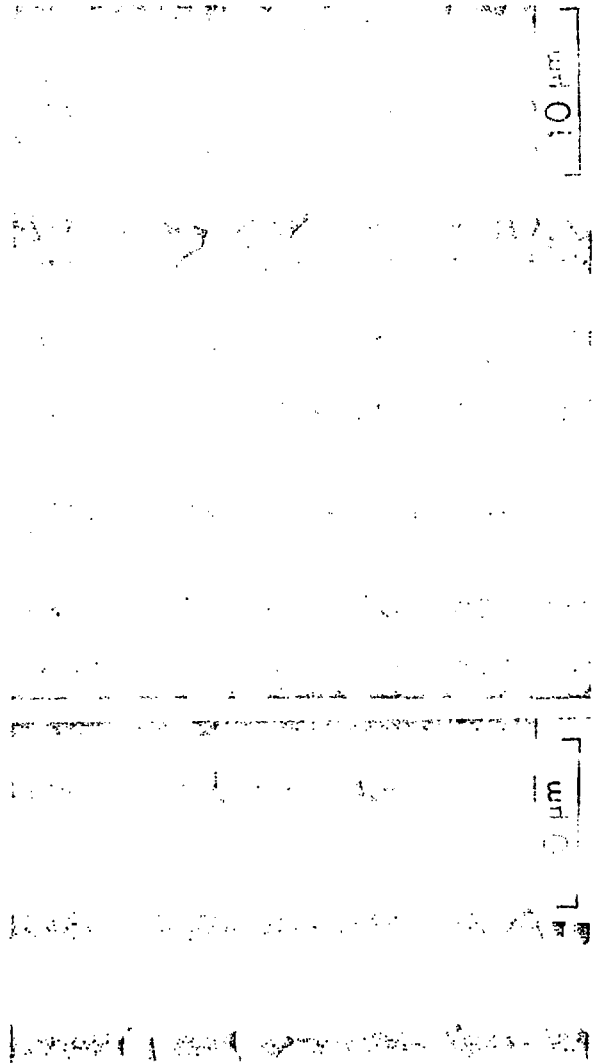


FIGURE 2

INSULATOR CERAMIC COATING no. 14 (52.3% SiO₂, 40.5% BaO, 7.2% Al₂O₃)
ON Nb-1% Zr IRRADIATED AT ROOM TEMP.

WITH 100keV He⁺

WITH 250keV He⁺

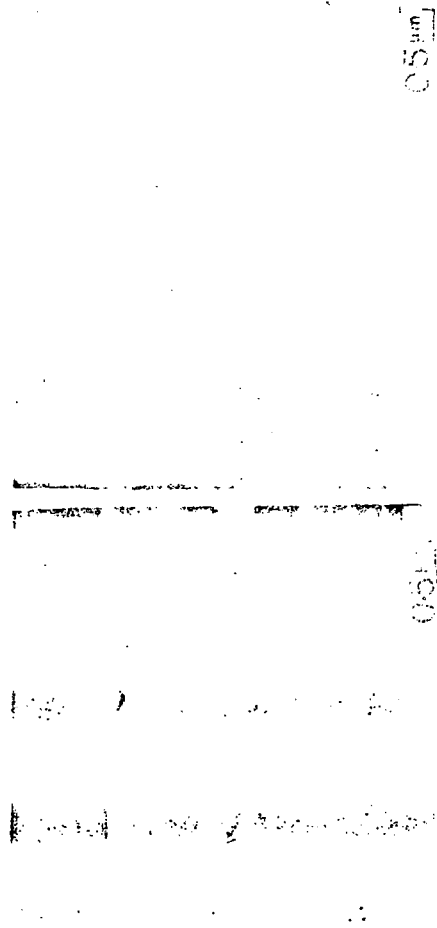


FIGURE 3

INSULATOR CERAMIC COATINGS IRRADIATED
300°C WITH 100keV He⁺

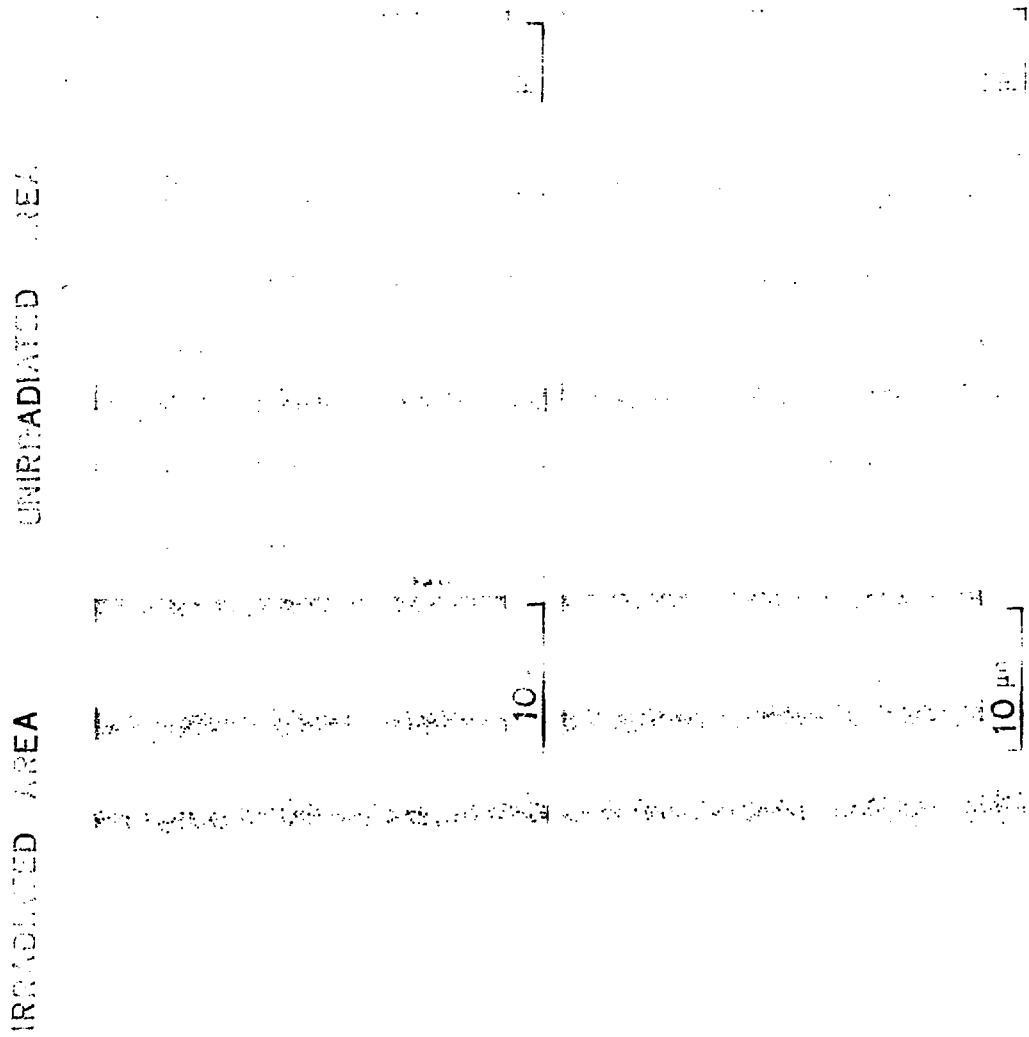


FIGURE 4