

Characterization and Modeling of Secondary Phase Evolution in an Irradiated Zr-1.0Nb Alloy

Matthew Swenson – University of Idaho – swenson@uidaho.edu

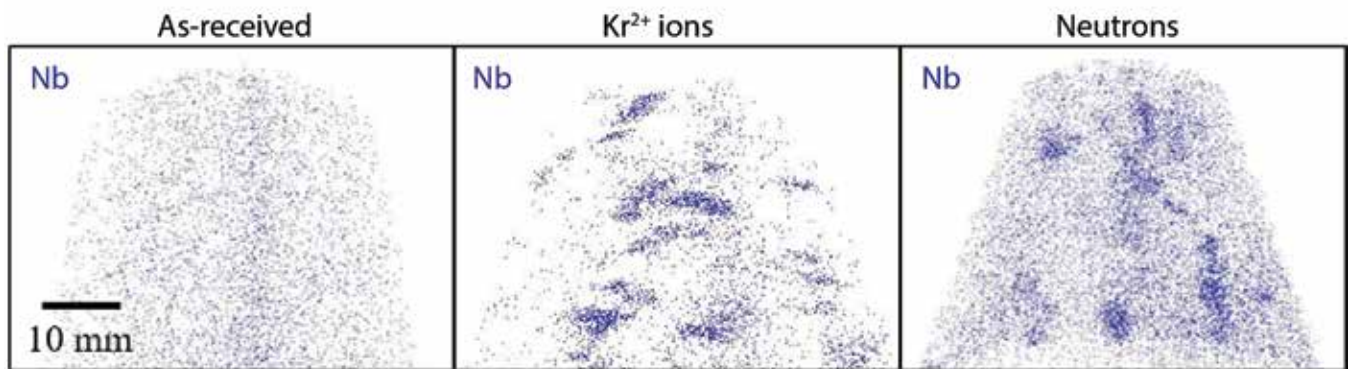


The objective of this study is to characterize the evolution of Nb-rich clustering and solute migration in a Zr-1.0%Nb alloy irradiated with either neutrons or Kr^{2+} ions to otherwise similar conditions (5 dpa at 310°C), enabling isolation of the dose-rate effects. Zirconium-based alloys are commonly used as cladding materials in existing reactor applications because of their low thermal-neutron absorption cross section, good corrosion resistance at high temperature, and high resistance to irradiation-induced swelling and creep. More recent development of Zr-based alloys with small amounts of niobium has also demonstrated improvement in corrosion resistance and the ability to minimize irradiation-induced linear growth. In this study, an RXA Zr-1.0%Nb alloy was irradiated with Kr^{2+} or neutrons to a common dose of ~ 5 dpa (each at 310°C). Atom-probe tomography is used to quantify any solute-cluster morphology and the solute concentrations in the matrix, clusters, and preexisting β -Nb precipitates before and after each irradiation. This approach enables systematic characterization of the Nb-solute migration as a result of irradiation, along with an evaluation of dose rate effects and the ability for Kr^{2+} ions to emulate neutron irradiation at low dose.

Results

In this study, atom-probe tomography (APT) is used to systematically quantify the migration of Nb solutes due to each irradiation. Prior to irradiation, only β -Nb precipitates are observed, with a surrounding matrix containing 0.59 at% Nb, which is close to the solubility limit for Nb [1,2]. Following both irradiations, nanoscale Nb-rich clusters are found within the matrix, consistent with prior observations [1,3,4], and are likely elongated or needle-shaped, with an average aspect ratio of $\sim 2:1$ measured using APT. The irradiation-induced clustering coincides with a reduction in matrix composition of Nb to 0.36 at% and 0.49 at% following Kr^{2+} ion or neutron irradiation, respectively. This evidence suggests the solutes migrate from the matrix to irradiation-induced clusters. The size of the Nb clusters are 4.51 ± 1.09 nm and 4.17 ± 2.23 nm following Kr^{2+} and neutron irradiation, respectively, while number density of the Nb clusters are $379 \times 10^{21} \text{ m}^{-3}$ and $253 \times 10^{21} \text{ m}^{-3}$ following the same irradiation conditions. Therefore, Kr^{2+} irradiations have resulted in higher volume fraction of Nb clustering at the same dose and temperature as neutron irradiation.

Irradiations to 5 dpa at 310 °C



Conclusion

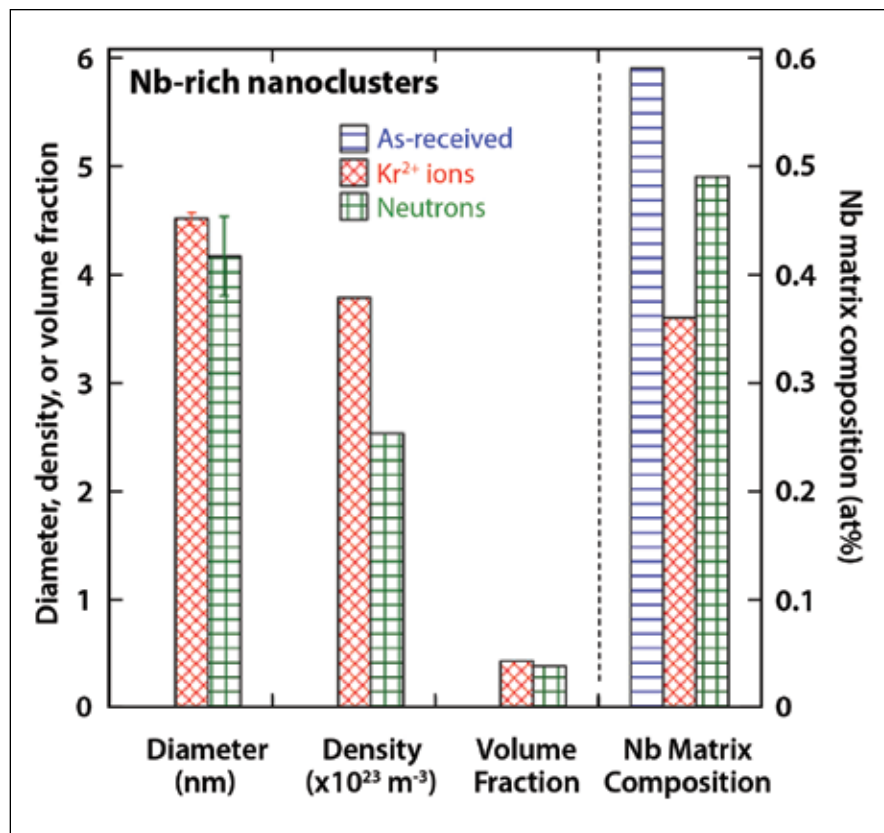
Irradiation with either Kr^{2+} ions or neutrons to 5 dpa at 310 °C have each led to the development of Nb-rich nanoclusters in a Zr-1.0%Nb alloy. Nanoclusters rich in Nb are expected to impede the growth of $\langle c \rangle$ loops in Zr-based alloys, likely reducing the effects of irradiation-induced linear growth [5,6]. The Nb clustering also lowers the amount of Nb in solid solution within the matrix, likely improving the corrosion resistance of the alloy [1]. The Kr^{2+} ion irradiation appears to be a reasonable emulation of neutron-irradiation effects on Nb solute migration, but some key differences are noted.

References

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Figure 1. Atom Probe concentration maps for Nb in Zr-1.0%Nb matrix. Following each irradiation, small Nb-rich nanoclusters are observed, with comparable size and number density.

Figure 2. Summary of Nb-rich cluster morphology follow each irradiation and the resulting evolution in the matrix composition of Nb in at%. Error bars for diameter represent the standard deviation of the mean.



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Publications

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 Evaluation of Nb-rich clustering in a Zr-1.0%Nb alloy following Kr²⁺ or neutron irradiation at 310 °C, Materials, Manuscript in preparation.

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University of Idaho	Matthew Hu (collaborator), Saheed Adisa (collaborator), Matthew Swenson (principal investigator)