§10. In-situ Optical Measurements of Decrystallization Induced by Ion Bombardment on Er₂O₃

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i) Introduction Optical methods are potentially useful for characterization of radiation-induced defects in ceramics blanket materials. Feasibility studies of the optical methods are undertaken for *in-situ* characterization of the radiation-induced defects in Er_2O_3 by ion bombardment. The characterization is important for qualification of Er_2O_3 coatings as electric insulation of Li/V-alloy blanket systems and as hydrogen permeation barriers.

A potentially useful luminescence band in 640-690 nm is identified as $4f^{11} {}^{4}F_{9/2} - {}^{4}I_{15/2}$ transition of Er^{3+} at C_{2} cation sites. It has been demonstrated that ion bombardment on a plasma splay coating sample ¹) and a sintered bulk sample ²) quenches preferentially the luminescence band.

ii) Modeling of luminescence quenching due to decrystallization by \mathbf{Ar}^+ ion bombardment For a given local displacement per ion (Φ), a displacement per atom (dpa: C) at a depth (x) and an ion fluence (f) is written as,

$$C(x;f) = \frac{1}{Y} \int_0^\gamma \Phi(x - \gamma') d\gamma' \quad (x \ge \gamma), \qquad (1)$$

where $\gamma = f \times Y/n$ is a surface erosion depth, and Y and n are a sputtering yield and an atomic number density of the target, respectively. Fractional number of emitters, *i. e.* Er^{3+} cations, may be depopulated by an amount of $\alpha \times C$, where α is a fraction of the dpa which contributes to depopulate the emitters. The depopulation of the emitting cations can be ascribed to decrystallization of the target. Figure 1 shows a measurement of luminescence quenching with Ar^+ (33 keV) ion fluences and fitting curves using the dpa given by Eq. 1. The local displacement per ion for $\mathrm{Er}_2\mathrm{O}_3$ was calculated by using the TRIM code. The fitting is significantly sensitive to sputtering yield data used in the dpa calculation.

iii) Luminescence quenching by \mathbf{H}^+ ion bombardment Figure 2 shows ion-induced luminescence spectra and its quenching with \mathbf{H}^+ (34 keV) ion fluences. Intensity of the luminescence band at 640 – 690 nm preferentially decreases as the ion fluence increases. The relative intensity decreases rapidly until the ion fluence attains to $2 \times 10^{21} / \mathrm{m}^2$. At larger ion fluences, however, the relative intensity appears to have a finite value independent on the ion fluence. During the \mathbf{H}^+ ion bombardment, not only decrystallization but also recovery may take place in the target at a finite temperature. In steady states, the luminescence intensity reaches at a value determined by rates of displacements and recombinations which may depend on temperature, projectile ion species and kinetic energies. This issue will be addressed in future studies.



Fig. 1: Luminescence quenching with Ar^+ (33 keV) ion fluences on a sintered Er_2O_3 sample ³⁾.



Fig. 2: Ion-induced luminescence spectra and its quenching with H^+ (34 keV) ion fluences on a sintered Er_2O_3 sample ⁴⁾.

- 1) T. Tanaka et al., J. Nucl. Mater. (2011) vol.417 794.
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- 4) D. Kato *et al.*, presented at the 29th JSPF annual meeting, Nov. 27-30, 2012, Kasuga, Japan.