Structural Characterization and Luminescence Properties of Er_xSc_{2-x}Si₂O₇ Prepared by RF Sputtering

A. Najar^{1*}, H. Omi^{1,2}, T. Tawara^{1,2}

¹NTT Basic Research Laboratories, NTT Corporation, 3-1, Morinosato-Wakamiya, Atsugi, Kanagawa 243-0198, Japan

²NTT Nanophotonics Center, NTT Corporation, 3-1, Morinosato-Wakamiya, Atsugi, Kanagawa 243-0198, Japan

E-mail: Adel.Najar@lab.ntt.co.jp

Er related luminescence at 1.53 μ m from Er-doped materials has been under intense investigation [1–2] for its wide applications in optical telecommunications and Si photonics. However, the optical gains and luminescence efficiencies are limited by the low doping concentration of Er due to its low solid solubility in most host materials. A promising solution is to use Er-RE compounds (RE (Y, Yb, Tm) with same radii than Er), which permit to insert gradually Er ions inside a proper crystalline structure by substituting RE ions with Er ions. However, Er-RE silicate with RE radii (example: Sc (0.75Å) smaller than Er (0.881 Å) have not been enough investigated.

In this work, Er-Sc disilicate thin films were grown by RF-sputtering by alternating 15 nm thick layers of Er₂O₃, Sc₂O₃ separated by 15 nm SiO₂ layer, than deposited on 50 nm of Er₂O₃. These layers were deposited on SiO₂ (1.3 µm)/c-Si (100) substrate. After annealing at 1250 °C, Er and Sc diffuse in the SiO₂ layers with concentration of 6.7×10^{21} and 1.4×10^{21} at/cm³, respectively. The synchrotron grazing incidence X-ray diffraction (GIXD) measurements coupled to selected area electron diffraction (SAED) analysis confirm the presence of Er_xSc_{2-x}Si₂O₇ compound. The dominant Er_xSc_{2-x}Si₂O₇ layer shows room-temperature photoluminescence peak at 1537 nm (Fig. 1 (a)). The shift peak emission compared to Y-Er silicate (where Y and Er have almost same ionic radii) is due to the small ionic radii of Sc³⁺ that enhanced the crystal field strength affecting the optical properties of Er³⁺ in Sc silicate phase. The effective excitation cross-section of Er in the formed compounds was equal to $\sigma_{ex} \approx 1.4 \times 10^{-21}$ cm² for measured lifetime of 38 µs at $\lambda_{ex} = 980$ nm. The Er-Sc silicate opens a promise way to prepare photonic crystal light emitting devices.



Fig. 1. (a) room temperature PL vs pump power for annealed sample at 1250 °C, inset cross-section TEM image after annealing with SAED image. (b) integrated PL intensity vs pump power, inset life time spectrum.

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