

Comparative study of crystal field analysis in Pr³⁺ and Yb³⁺-doped K₂LaX₅ (X= Cl⁻, Br⁻) ternary halides and Yb³⁺-doped A₃Lu(PO₄)₂ (A=Na⁺, Rb⁺) double phosphates. Charge transfer band observations of Yb³⁺-doped systems.

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The spectroscopic properties of praseodymium and ytterbium-doped ternary halides of K₂(La,Ln)X₅ (X=Cl⁻, Br⁻) type and ytterbium-doped double phosphates of formula A₃Lu(PO₄)₂ (A=an alkali metal: Na, Rb) were presented. Absorption, emission and luminescence excitation spectra were measured at 4, 10 and 293K in the wide spectral range (VUV-IR). Luminescence of the samples was excited using various pumping lines, including the synchrotron radiation. In the Pr³⁺ ion systems the Stark components determined from the absorption and emission spectra were simulated by using a phenomenological crystal field model for 4f² configuration of the Pr³⁺ ion in the K₂(La,Pr)X₅ crystals. The difference in the emission behaviour of the ternary chloride and bromide crystals were explained basing on C.F. effect and vibrational mode in the discussed systems. For the Yb³⁺-doped samples the strong fundamental ²F_{5/2} → ²F_{7/2} emission in the IR range as well as the CT emission bands were observed. The influence of the host lattice on charge transfer luminescence was investigated. The temperature effect and mechanisms of the observed phenomena are discussed.

Słowa kluczowe

Pr³⁺ ion, Yb³⁺ ion, Charge transfer emission, Crystal field calculation, Ternary halides, Double phosphates

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