

LUMINESCENCE OF NANO- AND MICROCRYSTALS EMBEDDED IN HALIDE MATRICES

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CsPbX₃ nanoparticles in CsX matrix (X=Cl, Br) on example of chlorine composite system





 $10^{\circ} \pm 10^{\circ} \pm 1$

Coherent effects Superluminescence



Excitation spectra







Ce³⁺ activated LaX₃ particles embedded into NaX hosts (X=Cl, Br, I)





Micrographs of NaX-LaX₃-Ce fresh cleaved surface obtained by: secondary electron registration mode (a, c), cathodoluminescence mode (b, d)

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Luminescence spectra



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Excitation mechanisms of Ce³⁺ luminescence





Decay kinetics



Smaller quantity of carrier trappings in microcrystalline phase (on example of LaCl₃:Ce microcrystals embedded into NaCl)



X-ray excited luminescence spectra



Excitation of Ce³⁺ centers by the reabsorption of the STE emission of a host



X-ray excited decay kinetics

Decay kinetics of STE of NaBr



Decay kinetics of Ce³⁺ centers in LaBr₃ microcrystals



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Eu²⁺ activated MeX₂ particles embedded into NaX (Me=Sr, Ba; X=Cl, I)





*NaCl-SrCl*₂(1 mol.%)-*EuCl*₃(0.02 mol.%)





*NaCl-BaCl*₂(1 *mol.%*)-*EuCl*₃(0.02 *mol.%*)





Micrographs of NaCl-MeCl₂-Eu freshly cleaved surface obtained by: secondary electron registration mode (a, c), cathodoluminescence mode (b, d)



Entering of Eu²⁺ ions in MeCl₂ microcrystals



Excitation mechanisms of Eu²⁺ luminescence



Decay kinetics



Effect of anion substitution on aggregation processes





The increase of excitonic luminescence intensity and shortening of decay time were revealed for CsPb(Cl, Br)₃ nanoparticles embedded into Cs(Cl, Br) host that could be explained by the coherent effects manifestation

The efficient formation of Ce³⁺ and Eu²⁺ doped lanthanum, strontium and barium halide microcrystals was shown into more stable natrium halide hosts. The luminescence properties are similar to the ones of corresponding single crystals. Smaller amount of carriers trapping was registered for embedded particles

The excitation mechanisms of activator luminescence were established for optical and X-ray excitations





Thank you for your attention!



