Taras Shevchenko National University of Kyiv
Institute of Physics NAS of Ukraine
Minor Academy of Sciences of Ukraine
Ukrainian Physical Society
under the auspices of
Representative Office of Polish Academy of Sciences in Kyiv

Dedicated to the 100th anniversary of the National Academy of Sciences of Ukraine

SPECTROSCOPY OF MOLECULES AND CRYSTALS

Book of Abstracts of XXIII Galyna Puchkovska International School-Seminar

(Kyiv, Ukraine, September 20–25, 2017)

Synthesis, Structure and Spectral-Fluorescent Properties of Organic-Inorganic Perovskite CH₃NH₃PbI₃ Films

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Organic-inorganic halide perovskites APbX₃ (A – CH₃NH₃, X – Cl, Br, I) attract now extensive scientific interest as a potent photoactive matrix for solar energy harvesting [1]. These compounds demonstrate high photovoltaic conversion efficiencies in laboratory conditions at relatively low costs of production and processing, having the only major drawback to overcome, viz. low durability.

Hitherto, there are no data in the literature about the effect of the CH₃NH₃I: PbI₂ ratio in the starting work solution on the structure and properties of organic-inorganic films of halide perovskites CH₃NH₃PbI₃. In this work, the microstructure, phase composition, and spectral-fluorescent properties of thin films of this organic-inorganic perovskite deposited on a glass substrate via spin-coating and heat-treated at 80 °C or 150 °C were studied at different reagent ratios in the initial solutions.

The shape and particle size of the films depend strongly on the reactants stoichiometric ratio. When the $CH_3NH_3I:PbI_2$ ratio is 1:1, the film consist of needle-like particles located along the substrate plane. For the ratio 2:1, the round-shaped particles were obtained. With the further increase of methyl ammonium iodide content (3:1), a smooth transition from round to polyhedral particles was observed. In all three cases, the $CH_3NH_3PbI_3$ particles are predominantly located in a single layer.

It is found that as the CH₃NH₃I content increases, the UV/Vis absorption spectra become more selective – the intensity increases in the region of 350-400 nm while the absorption in the visible spectrum range decreases. In the initial solution in DMF, the absorption range does not exceed 350 nm, with the most long-wavelength band at 327 nm indicating partial complexation of the mixture components in solution. Relative fluorescence intensity of the studied CH₃NH₃PbI₃ films increases with the higher content of CH₃NH₃I in the initial solution. This regularity is observed for films after heat treatment at both 80 °C and 150 °C, although in the latter case the fluorescence is abated substantially, probably because of partial destruction of perovskite [1].

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