The comparison of photoluminescence decay in YAG:Er, ZnO and SiO₂ crystals

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We report on the intensity and time resolved photoluminescence (PL) measurements in the visible spectral range at wavelengths 350-800 nm using the phase delay method under sine wave UV excitation. Yttrium aluminium garnet (Y₃Al₅O₁₂, YAG), zinc oxide (ZnO) and silicon oxide (SiO₂) are crystalline materials, known for their excellent optical properties and mechanical, chemical and temperature stability. The YAG:Er crystals were grown by the micro-pullingdown method at the Institute of Physics in Prague [1], ZnO crystals by hydrothermal growth at the Institute of Chemistry in Bratislava [2] and the SiO₂ micro powder was purchased from Sigma-Aldrich and expoosed to inductively coupled plasma (ICP) at the Institute of Physics to create surface-related defects. Their PL spectra were measured at a room temperature using the UV LED sine wave modulated by 50 MHz Keithley 3390 generator, dispersive monochromator, 10 MHz red-enhanced photomultiplier, 1 MHz current amplifier 10⁵ V/A, 200 MHz oscilloscope, and a 100 kHz lock-in amplifier. The sensitivity of the oscilloscope is about 3 orders of magnitude lower than the sensitivity of the lock-in amplifier. Our setup allows to measure the mean PL decay with a time resolution of about 10 ns at 100 kHz. While YAG:Er shows well resolved PL peaks with mean decay time of several μ s related to the Stark splitting of Er³⁺ (4f¹¹) states, the defect-related PL spectra of ZnO and SiO₂ show broad defect-related bands with significantly faster mean decay times (in the order of tens or hundreds ns in ZnO, depending on doping, and below 10 ns in SiO₂).

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