

Red-Emitting Lu₂S₃:Ce Scintillator Prepared by Micro-Pulling-Down Crystal Growth

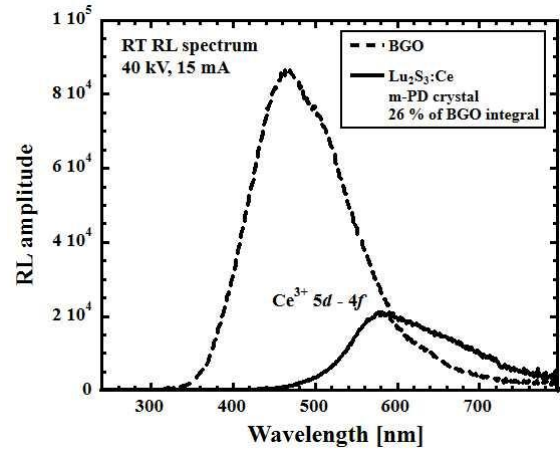
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In recent years, scintillators emitting in the red spectral region have attracted increasing attention due to the rapid development of semiconductor-based photodetectors, whose sensitivity to red photons surpasses that of conventional photomultiplier tubes. Among potential red-emitting scintillator materials, sesquisulfides represent a largely unexplored class. Recent results demonstrated the potential of sulfide-based crystalline materials: KLuS₂:Pr³⁺ crystalline plates are excellent candidates for time-of-flight positron emission tomography, exhibiting an ultrafast scintillation decay time (< 1 ns) and a scintillation light yield of approx. 7000 ph/MeV (1), red-emitting Lu₂S₃:Ce, grown by the chemical vapor transport method, has been reported in Ref. (2). To the best of our knowledge, bulk sesquisulfide crystals with dimensions exceeding approx. 1 mm have not yet been reported. Sulfide materials have previously been studied mainly as novel laser media, while their scintillation properties remain insufficiently studied.



Room temperature radioluminescence spectrum of Lu₂S₃:Ce grown by micro-pulling-down method in comparison with radioluminescence spectrum of BGO.

In this contribution, we present a systematic investigation of the cerium-doped sesquisulfide Lu₂S₃. This material benefits from the high atomic number of lutetium ($Z = 71$) and a relatively high compound density ($\rho = 6.25 \text{ g}\cdot\text{cm}^{-3}$). The resulting high density and effective atomic number of Lu₂S₃:Ce ($Z_{\text{eff}} = 66.8$) are highly desirable properties for scintillators, as they enhance radiation detection efficiency and enable the fabrication of compact detectors compared to conventional materials with lower density and Z_{eff} .

The samples were prepared using the micro-pulling-down method, which allows rapid laboratory-scale growth of high-quality single crystals with dimensions of several millimeters. Phase purity and structural quality were verified by X-ray diffraction. Lu₂S₃:Ce exhibits a broad emission band centered at 580 nm, attributed to the allowed 5d–4f transition of Ce³⁺ ions. The integrated intensity is approximately 26 % of that of Bi₄Ge₃O₁₂ (BGO) single crystal. No other optically active impurities were detected. Owing to the dipole-allowed nature of the Ce³⁺ transition, the material shows a fast scintillation decay time on the order of 30 ns. Photoluminescence and scintillation characteristics of a series of micro-pulling down-grown samples will be presented. The light yield was evaluated using pulse-height spectroscopy, and the presence of possible slow decay components will be discussed.

(1) V. Jarý et al., “Efficient Ultrafast Scintillation of KLuS₂:Pr³⁺ Phosphor: A Candidate for Fast-Timing Applications,” *Phys. Rev. Applied* **19** (2023) 034092

(2) J. C. van’t Spijker et al., “Lu₂S₃:Ce³⁺, A new red luminescing scintillator,” *Nucl. Instr. and Meth. in Phys. Res. B* **134** (1998) 304-309

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