

# YAG:Ce3+/4+ aerogels: efficiency, timing, diffusion and self-absorption

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### Oral Sessions | Tuesday, July 9<sup>th</sup> 2024

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#### T-NA2 YAG:Ce<sup>3+/4+</sup> aerogels: efficiency, timing, diffusion and self-absorption

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Detection of radionuclides that decay directly to the ground state in the atmosphere requires to detect the beta or alpha radiation directly, which is very challenging due to their short mean free path in air at normal pressure. This suggests that the highest detection efficiency is obtained when the radionuclide is incorporated in the sensing part of the detector, which is applied with different gas radionuclides in ionisation chambers (gas-gas mixing) and for liquid solution in the liquid scintillation counting (LSC) technique (gas-liquid mixing). We hereby introduce the concept of gas-solid mixing to design an efficient detector, wherein the radioactive gas is introduced into a highly porous transparent inorganic scintillator. In this frame we have prepared YAG:Ce aerogels from sol-gel nanoparticle dispersion.

Post-synthesis thermal treatment (sintering/calcination) at several temperatures was applied to the aerogels demonstrating a significant impact on the cerium oxidation state and transparency (figure 1). The focus of this work is also the investigation of the aerogel optical properties after being altered by sub-calcination/sintering temperature treatments in H<sub>2</sub> or air ambient gases. This low-temperature (400°C) treatment has managed to significantly change the ratio of  $Ce^{3+/4+}$  population in the sample, thus introducing a novel, easy and reversible approach to change scintillating properties of YAG:Ce aerogel.

In this presentation, the effect of the thermal treatment on the transparency, diffusion, self-absorption, photoluminescence and scintillation efficiency and timing will be discussed in the frame of their use as sensor in the Triple to Double Ratio method, the standard technique used for liquid scintillation and the metrology of radionuclides.



Figure 1: YAG:Ce aerogels under various thermal treatments

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In recent years cesium lead halide perovskite nanocrystals have attracted attention in many application fields, among which that of radiation detection. Thanks to quantum confinement effect they feature ultrafast scintillation decays [1]. Therefore, they have been identified as promising fast emitters in scintillating metamaterials intended to build up novel detectors requiring fast timing [2]. Such detectors are particularly needed in the time-of-flight (TOF) techniques in medical imaging as well as high energy physics. However, the practical usage of nanocrystals requires dealing with two major issues. Poor chemical stability can be overcome by their incorporation in a suitable, e.g. polymer, matrix to form nanocomposite. Low stopping power calls for fabrication of nanocomposites with the highest possible nanocrystalline loading keeping their reasonable level of transparency.

In this work we study synthesis, structural and optical characteristics including the timing performance of cesium lead bromide and cesium lead bromo-chloride nanocrystals embedded in the polystyrene matrix. We focus on synthesis routes for nanocomposites with high nanocrystalline loading up to 40%. However, with higher loading favourable luminescent properties of nanocrystals can be compromised by their agglomeration leading to reduced transparency and consequently the reduced light extraction from prepared nanocomposites. To limit this effect, we use advanced polymerization techniques and copolymerizable ligands. The level of nanoparticle agglomeration is studied by confocal microscopy. We also study scintillation characteristics of prepared nanocomposites, especially scintillation decays. We assess the nanocomposite time resolution using a setup described in [3]. Prepared nanocomposites feature significantly better time resolution with respect to the currently used standalone conventional scitulilators, such as LYSO:Ce crystal.

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