Subpicosecond phenomena in scintillators for fast timing in future CERN experiments and medical applications

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COLLABORATION



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BACKGROUND

Study of scintillation crystals





With Dr. Stanislay Burachas



Positron emission tomograph

BGO $(Bi_4Ge_3O_{12})$

PWO (PbWO₄)

ALICE experiment

The results obtained form 1995 to 2009 are published in 20 papers in journals like *J. Cryst. Growth, NIMA, etc.*



CURRENT MOTIVATION

FAST TIMING

TARGET





CRYSTAL CLEAR COLLABORATION



STATE OF THE ART





TWO ROADS TO 10 ps RESOLUTION



THE ROADS ARE PAVED BY EXPERIMENTAL TECHNIQUES



Two-photon absorption

Probe for

free carrier

absorption

THE ROADS ARE PAVED BY EXPERIMENTAL TECHNIQUES

Time-resolved photoluminescence spectroscopy

EXCITATION:

Yb:KGW oscillator (*Light Conversion Ltd.*) emitting at 1030 nm. **80 fs** pulses at 76 MHz repetition rate



Harmonics generator (HIRO, *Light Conversion Ltd*.); the third **343 nm (3.64 eV)** and fourth **254 nm (4.9 eV)** harmonics



THE ROADS ARE PAVED BY EXPERIMENTAL TECHNIQUES

Time-resolved photoluminescence spectroscopy

800 750 700 650 600 550 500 450 400 350 300 250 [nm

DETECTION:



SCINTILLATORS UNDER CURRENT STUDY



Self-activated

gadolinium aluminum gallium garnet (GAGG, $Gd_3Al_2Ga_3O_{12}$)

PWO (PbWO₄)

Results on TRPL, PWO



Decay times: τ_1 = 3.8 ps and τ_2 = 683 ps @ 343 nm exc. τ_1 = 5.9 ps and τ_2 = 824 ps @ 254 nm exc.

Both at 343 nm and 254 nm excitation, the kinetics of luminescence within 400-500 nm and 500-600 nm are identical



Results on TRPL, PWO



Initial stage of spectrally integrated photoluminescence kinetics (dots), the instrumental response function (green line), and the best fit obtained using a bi-exponential decay function (red solid line)

Results on TRPL, GAGG:Ce



Luminescence spectra at two excitation wavelengths

At the same excitation pulse energy and absorption coefficient , PL intensity is considerably lower after predominant excitation of Gd^{3+} ions \Rightarrow nonradiative recombination during the excitation transfer to radiative recombination centers is important

Results on TRPL, GAGG:Ce

Luminescence kinetics at two excitation wavelengths



The major part of the luminescence follows the leading edge of the instrumental function. We observe a rise component with the rise time of 8 ns @ 254 nm exc. 2.5 ns @ 343 nm exc.

This is in consistence with the 2 ns rise time observed in GAGG:Ce under gamma irradiation [M.T. Lucchini et al., NIMA, **816**, 176 (2016)].

Results on two-photon absorption

Results on two-photon absorption, PWO



Results on two-photon absorption, PWO

Influence of gamma irradiation



Spectra of differential optical transmittance induced by 500 mJ/cm² pump at 395 nm Polarization: along and at 75[°] to the crystal axis b <u>Under</u> and <u>without gamma irradiation</u> (⁵⁷Co source of ionizing radiation (122 keV, 2 mCi) mounted at a distance of 1 cm from the plate surface)

Results on two-photon absorption, GAGG:Ce

Two-photon absorption in GAGG:Ce

pump 3.44 eV 16 pump 3.44 eV 0.19 mJ/cm^2 eV 10 0.19 mJ/cm² 14 - 3.34865 3.25197 12 DT signal (mOD) 3.15267 DT signal (mOD) -0.99495 - 3.05172 -0.19493 10 0.00515 0.20515 8 0.40516 .6 ps 0.6052 6 0.80521 1.00525 2.0054 -21/ 2.9 0.1 3.2 3.4 3.0 3.1 3.3 -2 -1 0 2 3 Δ Photon energy (eV) Delay (ps) Distortion caused by scattered laser beam Laser ~150 fs

Pump at 3.44 eV (360 nm)

Results on free carrier absorption

Results on free carrier absorption, PWO





Kinetics of optical density induced by short pulse excitation of PWO crystal at 254 nm, probed at 1030 nm, for different excitation pulse energy densities (indicated)

Initial part of the kinetics

Induced optical density at 1030 nm probe wavelength at 0.3 ps delay after short pump pulse at 254 nm versus pump pulse energy density

Results on free carrier absorption, YAGG:Ce



Kinetics of optical density induced by short pulse excitation of YAGG:Ce crystal at 254 nm, probed at 1030 nm, for different excitation pulse energy densities (indicated)



Initial part of the kinetics

Results on free carrier absorption, GAGG:Ce



Kinetics of optical density induced by short pulse excitation of GAGG:Ce crystal at 254 nm for different excitation pulse energy densities (indicated)



Initial part of the kinetics

The rise of FCA in GAGG:Ce, takes a few picoseconds, what is caused by delocalization of holes from Gd³⁺ to valence band.

This observation is consistent with the conclusion reported in [P. Dorenbos, ECS J. Solid State Sci. Technol. 2, R3001 (2013)] that the ground state of the ${}^{8}S-{}^{6}D_{7/2,9/2}$ intracenter transition of Gd³⁺ ions is located within the valence band.

Thank you for your attention