

## Solid state detectors

Si, Ge  
+ good energy resolution,  
~0.1 %  
- slow  
- low stopping power

## Scintillators

- bad energy resolution,  
~ 6 % as best  
+ fast – slow  
+ high – low stopping power

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## *wide bandgap semiconductors*

CdTe, CdZnTe, HgI<sub>2</sub>  
+ high stopping power  
- energy resolution ~ 1 %

## *new materials*

+ high light yield  
+ energy resolution ~ 3 %

	$E_{\text{gap}}$ [eV]	$E_{\text{e-h}}$ [eV]
Si	1.12	3.61
Ge	0.74	2.98
CdTe	1.47	4.43
HgI <sub>2</sub>	2.13	4.2
NaI(Tl)	5.9	15.3
CsI(Tl)	6.5	15.2

conduction band



Activator  
excited states



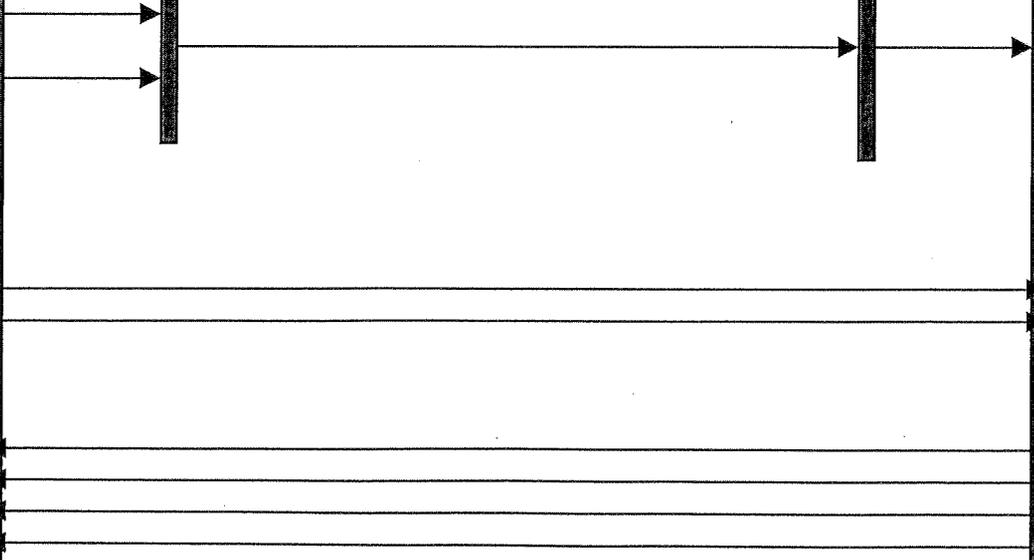
Activator  
ground state



valence band



band  
gap



## Inorganic scintillators

Scintillation photons are emitted from the activator excited states, fast transition, visible light.

Competing processes:

Electron arrives to excited states from which transition to ground state is forbidden. Additional small energy is required to lift the electron to other excited states (thermal excitation) with higher transition probability. This results in *phosphorescence radiation* radiation, or “*afterglow*”, which is a slow time component.

Some excited states cause radiationless transitions, which represents loss mechanism in the conversion of the radiation energy to scintillation light. This is the *quenching effect*.

	$\tau$ [ns]	$\lambda_{\max}$ [nm]	light yield [photons/MeV]
NaI(Tl)	230	415	44 000
CsI(Tl)	1000	550	60 000
BaF <sub>2</sub>	0.6/630	220/310	2200/7000
BGO Bi <sub>4</sub> Ge <sub>3</sub> O <sub>12</sub>	300	480	8000

## Scintillation process

5

- conversion to e-h pairs
- transfer to luminescence centre
- luminescence, i.e. photon emission

Number of emitted scintillation photons  $N_{ph}$  following absorption of gamma energy  $E_\gamma$

$$N_{ph} = \frac{E_\gamma}{\beta E_{gap}} SQ$$

$E_{e-h} = \beta E_{gap}$  - energy required to produce one e-h pair

$\beta \sim 2-3$  (mostly 2.3)

$S$  – transfer efficiency of the e-h pair to luminescence centre

$Q$  – efficiency for photon emission of the luminescence centre

Energy gap (eV)

NaI (Tl)	5.9
CsI(Tl)	6.4
BaF <sub>2</sub>	
fast	18.0
slow	10.6
ZnSe (Te)	2.6

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Energy required to produce one e-h pair

$$E_{e-h} = \beta E_{\text{gap}} \quad \beta \approx 2.3 \text{ (2-3)}$$

CsI (Tl)      assume  $E_{\gamma} = 1 \text{ MeV}$

No. of light photons

$$1 \text{ MeV} / 2.3 \times E_{\text{gap}} = 1\,000\,000 \text{ eV} / 2.3 \times 6.4 \approx 67\,000 \text{ photons}$$

compare with measured 60 000 photons

For other crystals

$$E_{\gamma}=1 \text{ MeV}$$

NaI (Tl)

$1\,000\,000 \text{ eV} / 2.3 \times 5.9 \text{ eV} \approx 73\,000 \text{ photons}$   
measured 40 000 photons

BaF<sub>2</sub>

fast  $1 \text{ MeV} / 2.3 \times 18.0 \text{ eV} \approx 24\,000 \text{ photons}$   
measured 1600 photons

slow  $1 \text{ MeV} / 2.3 \times 10.6 \text{ eV} \approx 41\,000 \text{ photons}$   
measured 8000 photons

## Radiative decay time for electric dipole emission

$$\tau_r \sim \lambda^2$$

more detailed

$$\tau_r = 1.5 \times 10^{-5} \frac{\lambda^2}{f^{1/9} (n^2 + 1)^2 n}$$

$f$  - oscillator strength of the transition

$n$  - refractive index

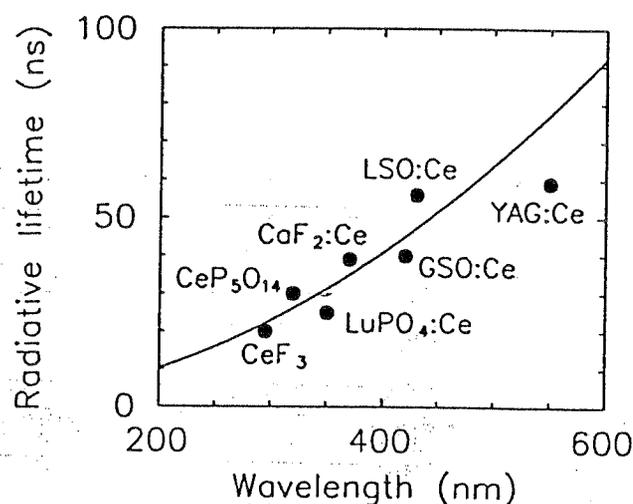


Fig. 2. The radiative lifetime vs wavelength. The oscillator strength,  $f$ , and refractive index,  $n$ , were taken equal to 0.0013 and 1.7, respectively. The points correspond to Ce d-f emission in different materials, as indicated. The decay times and emission wavelengths were taken: for GSO:Ce-Gd<sub>2</sub>(SiO<sub>4</sub>)O:Ce from [28], for LSO:Ce-Lu<sub>2</sub>(SiO<sub>4</sub>)O:Ce from [28], for YAG:Ce-Y<sub>3</sub>Al<sub>5</sub>O<sub>12</sub>:Ce from [29], for CaF<sub>2</sub>:Ce from [29], for CeF<sub>3</sub> from [3,6], for CeP<sub>5</sub>O<sub>14</sub> from [4].

High light yield  $\rightarrow$  low energy gap crystals  $\rightarrow$  long wavelength of the scintillation light

Fast crystals  $\rightarrow$  short  $\lambda$  (UV region)  $\rightarrow$  high energy gap

## Energy resolution

$$\left(\frac{\Delta E}{E}\right)^2 = (\delta_{sc})^2 + (\delta_{st})^2 + (\delta_n)^2$$

$\delta_{sc}$  – intrinsic resolution of the crystal

$\delta_{st}$  – statistical contribution

$\delta_n$  – dark noise contribution

For PM-tubes

$$\delta_{st} = 2.35x \left(\frac{1}{N}\right)^{\frac{1}{2}} x (1 + \varepsilon)^{\frac{1}{2}}$$

$N$  – number of photoelectrons

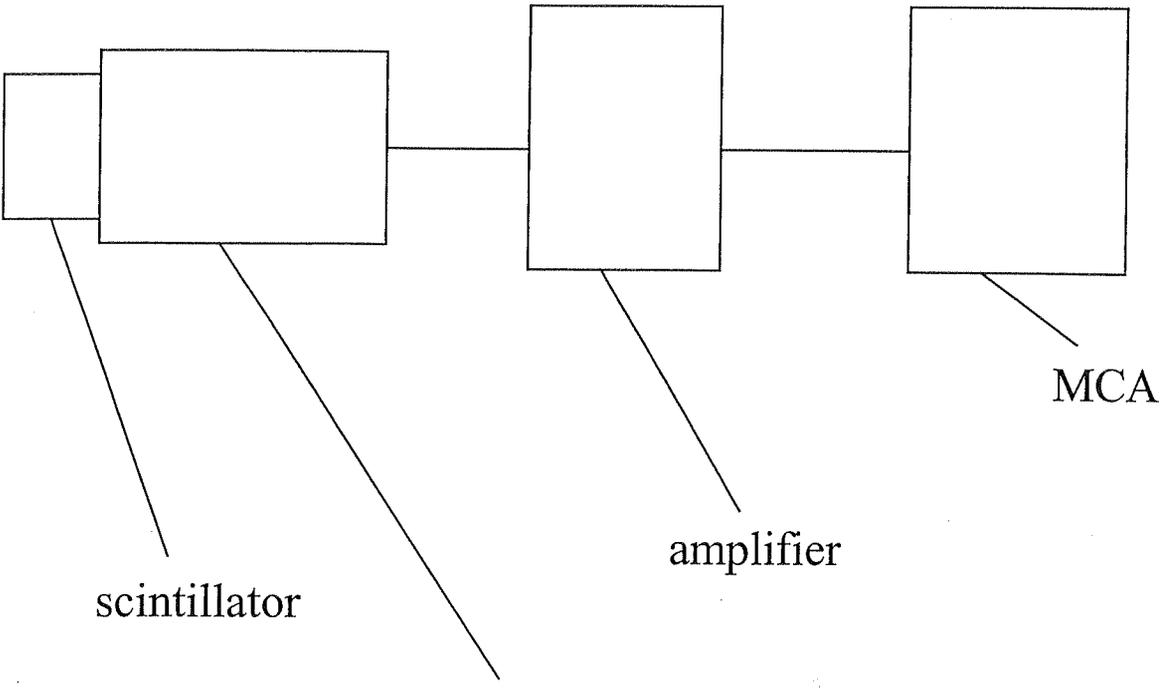
$\varepsilon$  – variance of the electron multiplier gain

For avalanche photodiodes (APD)

$$\delta_{st} = 2.355x \left(\frac{F}{N_{e-h}}\right)^{\frac{1}{2}}$$

$N_{e-h}$  – number of e-h pairs

$F$  – excess noise factor, reflecting the statistical fluctuation of the APD gain



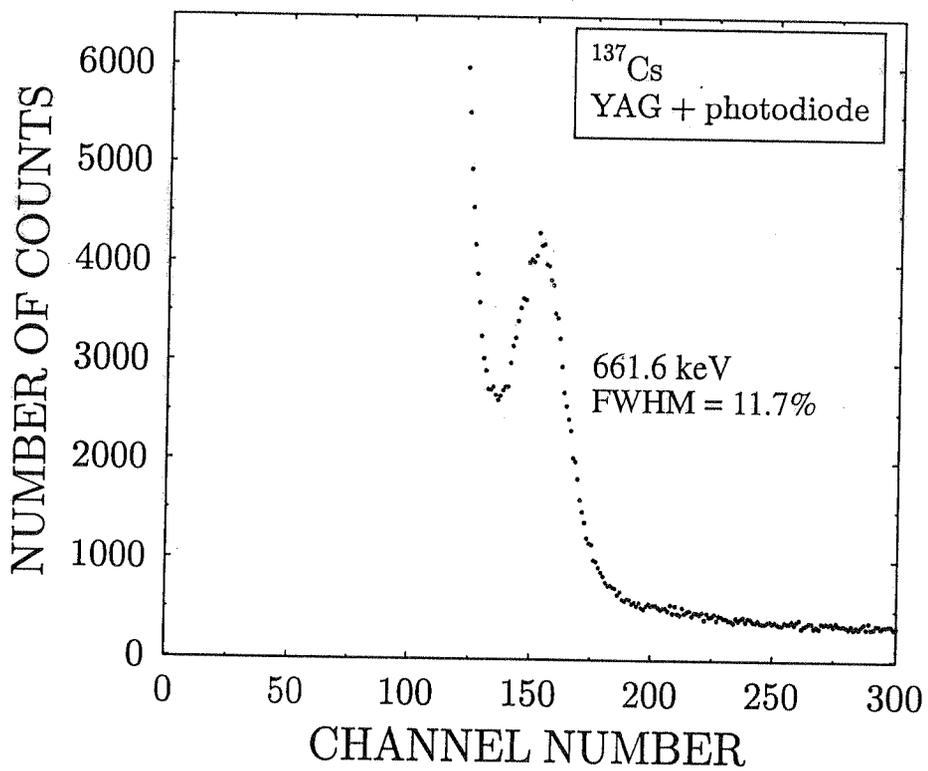
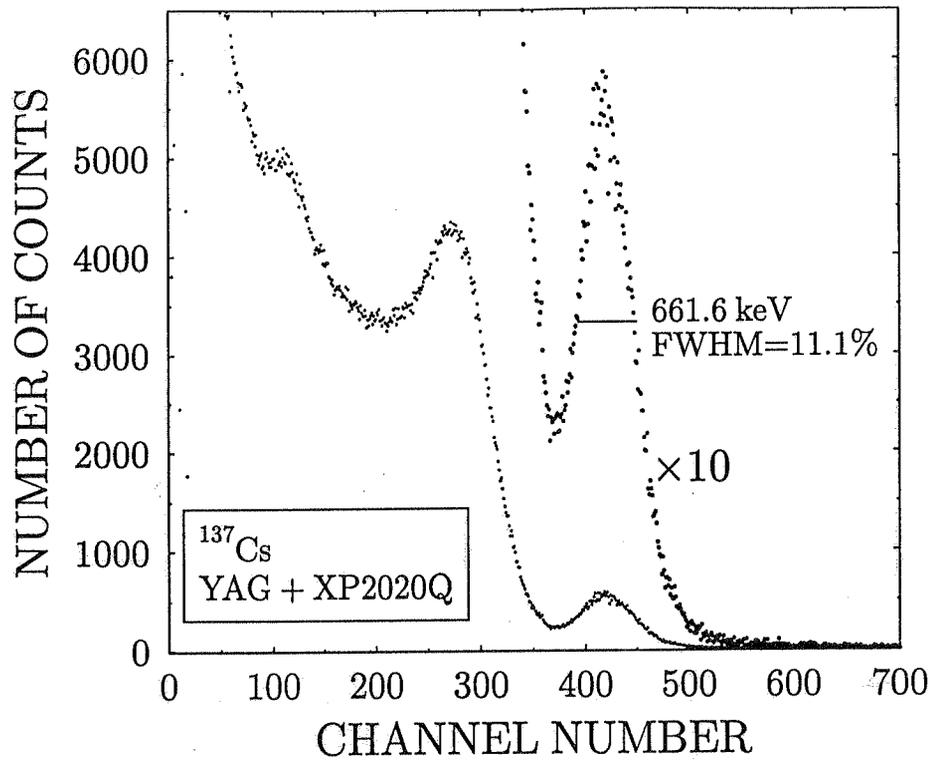
scintillator

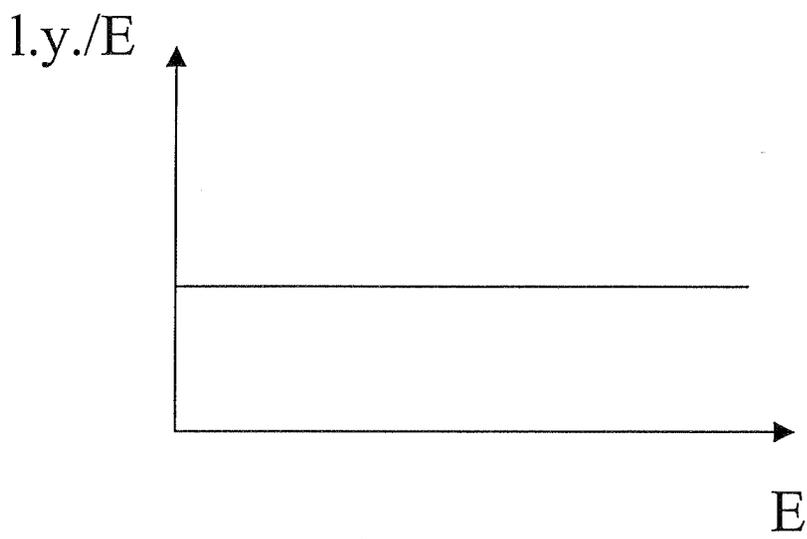
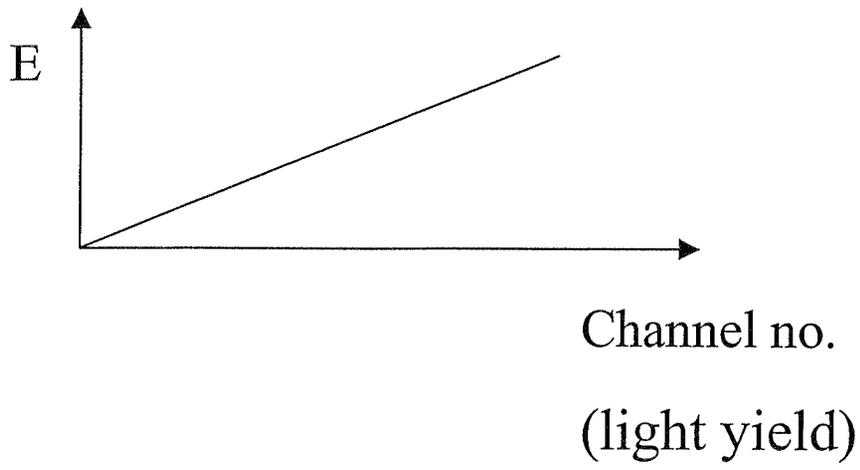
photodetector:

amplifier

MCA

- PIN photodiode
- PM-tube
- avalanche photodiode





# light yield nonproportionality in scintillators

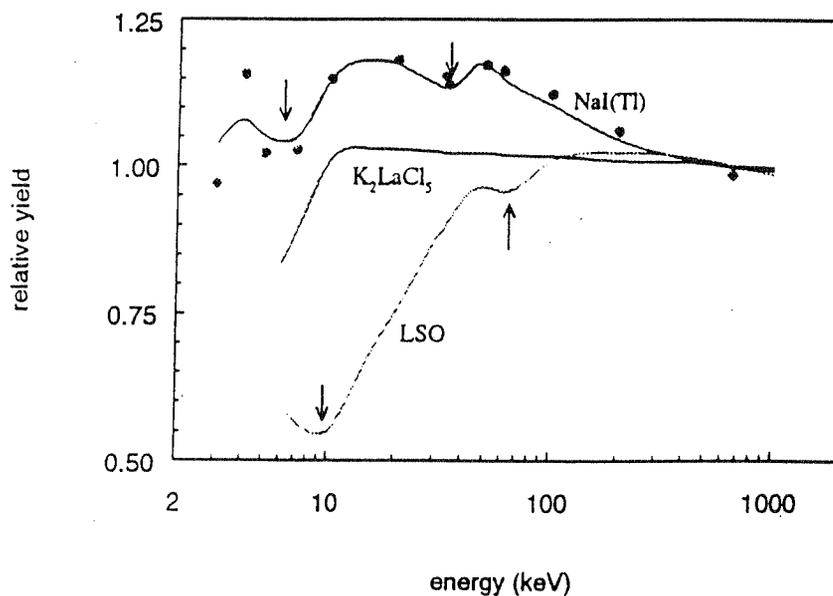


Figure 1: Response curves of NaI(Tl<sup>+</sup>), Lu<sub>2</sub>SiO<sub>5</sub>(Ce<sup>3+</sup>)(LSO), and K<sub>2</sub>LaCl<sub>5</sub>(Ce<sup>3+</sup>), normalized to unity at  $E_\gamma=662$  keV. The experimental data points, not shown in the Figure, for NaI(Tl<sup>+</sup>) can be found in [3]. For K<sub>2</sub>LaCl<sub>5</sub>(Ce<sup>3+</sup>) and Lu<sub>2</sub>SiO<sub>5</sub>(Ce<sup>3+</sup>) see [5]. The data points are results from Monte Carlo simulations of the scintillation process in NaI(Tl<sup>+</sup>). The arrows show the position of K-shell and L-shell binding energies.

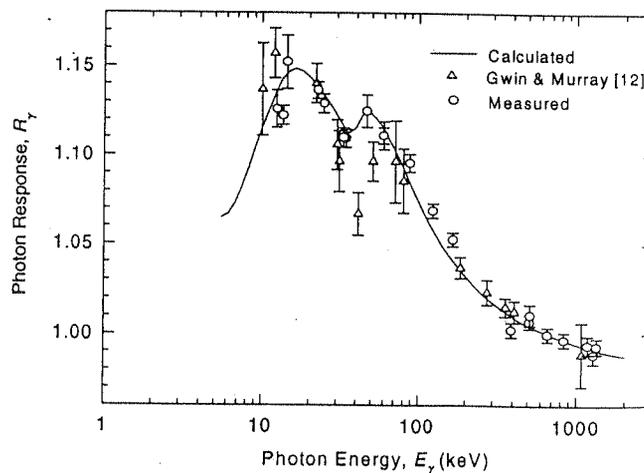
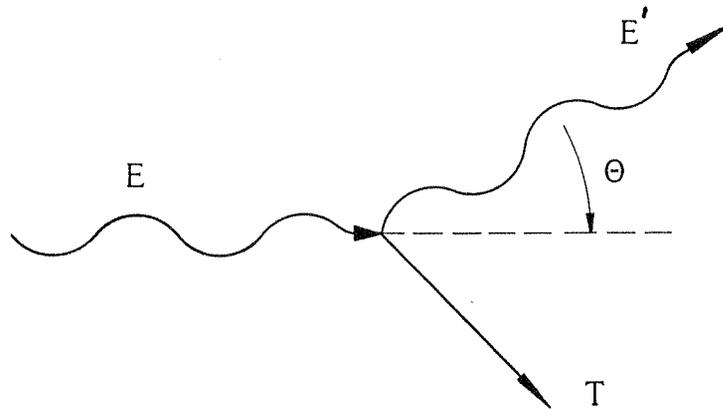


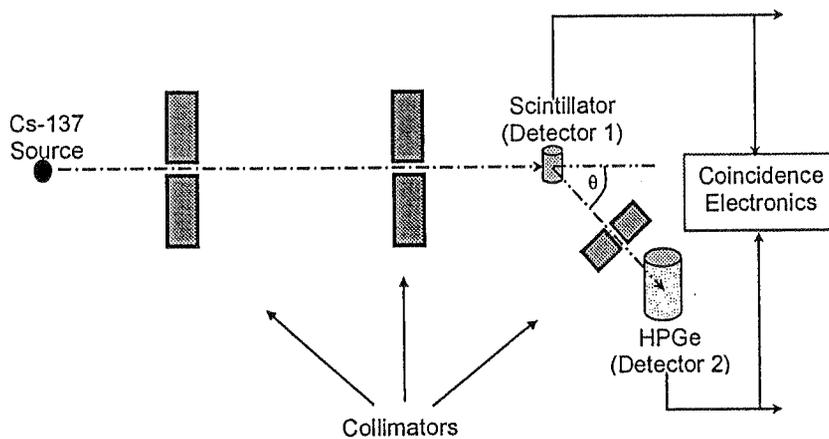
Figure 4: CsI(Tl) calculated and measured photon responses  $R_\gamma (=L_\gamma/E_\gamma)$ . Results from Gwin and Murray [12] have also been included. Data are normalized to unity at 661.66 keV.

# Compton scattering



$$E' = \frac{E}{1 + (1 - \cos \theta) \frac{E}{0.511}}$$

$$T = \frac{E^2(1 - \cos \theta)}{E(1 - \cos \theta) + 0.511}$$



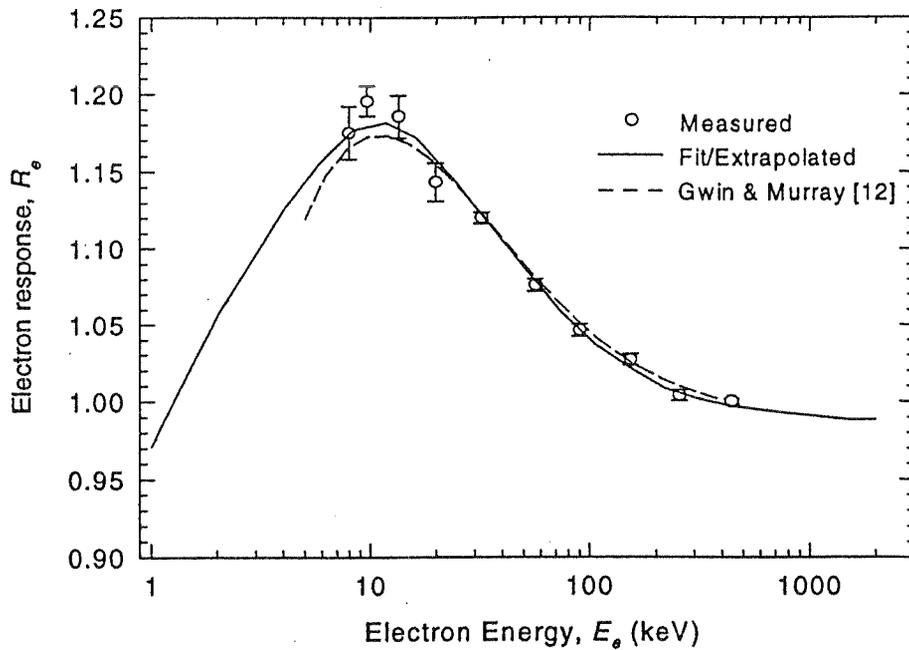


Figure 3: CsI(Tl) electron response  $R_e(=L/E_e)$  measured using the CCT along with a smooth fit and low and high energy extrapolations used in photon response calculations. Also shown are previously reported calculated results from Gwin and Murray [12]. Data are normalized to unity at 442 keV.

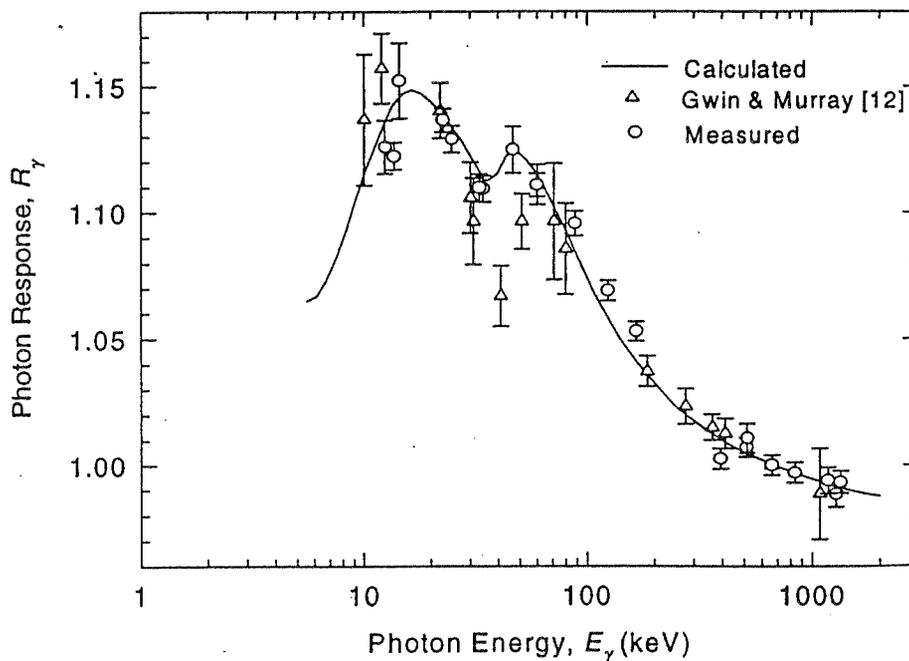


Figure 4: CsI(Tl) calculated and measured photon responses  $R_\gamma(=L_\gamma/E_\gamma)$ . Results from Gwin and Murray [12] have also been included. Data are normalized to unity at 661.66 keV.

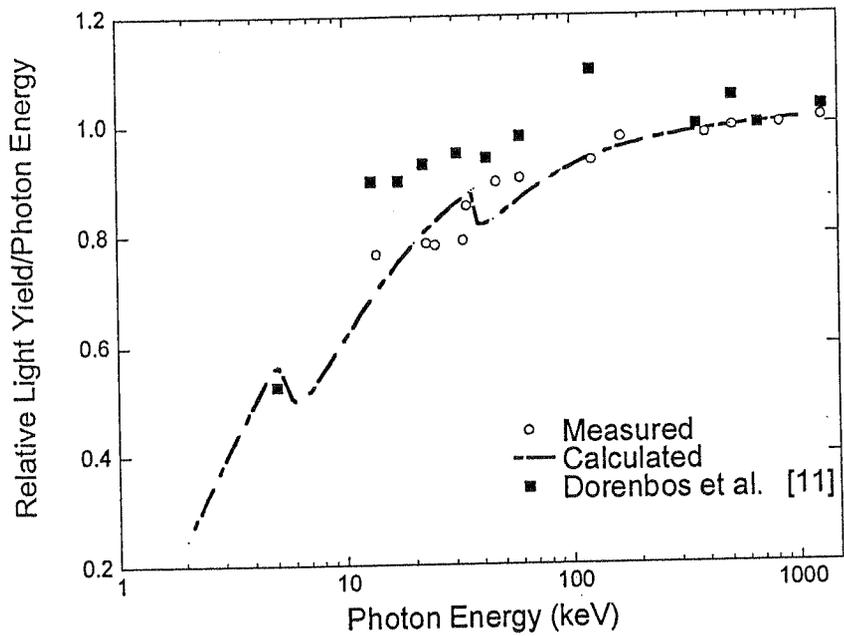


Fig. 5. BaF<sub>2</sub> calculated and measured photon responses. A smooth line is drawn through the calculated response to guide the eye.

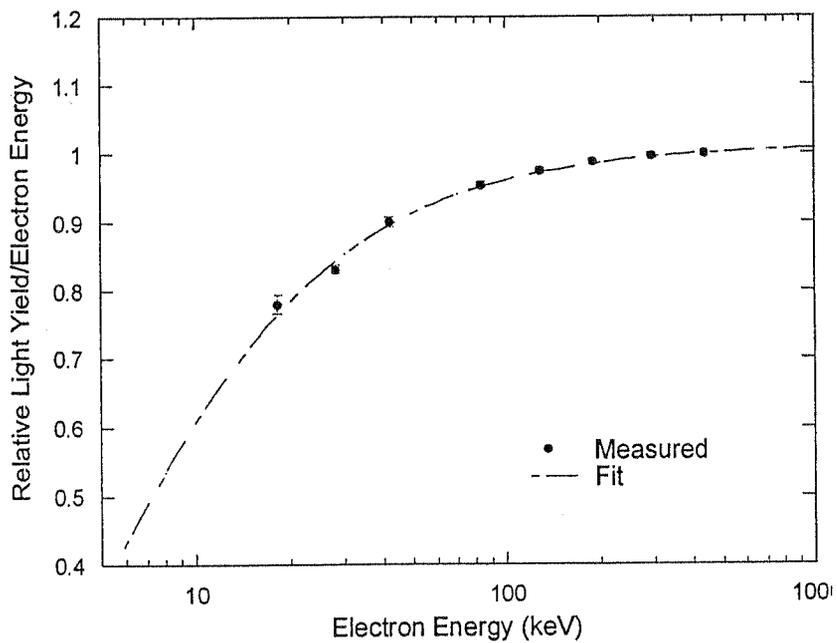


Fig. 4. BaF<sub>2</sub> measured electron response. The response increases by 22% as the electron energy increased from 18 keV to 436 keV.

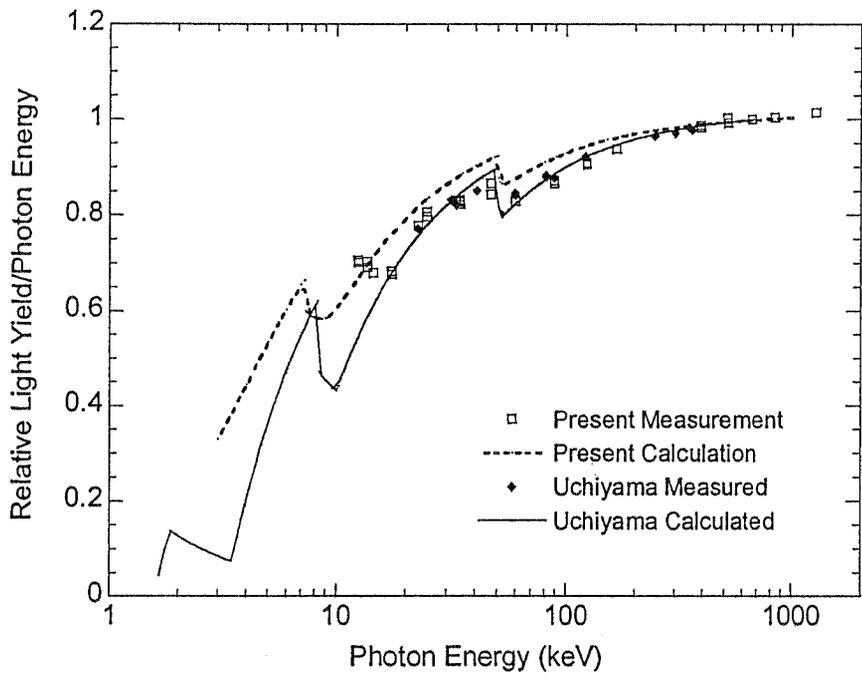


Fig. 3. GSO calculated and measured photon responses. A smooth line is drawn through the calculated response to guide the eye.

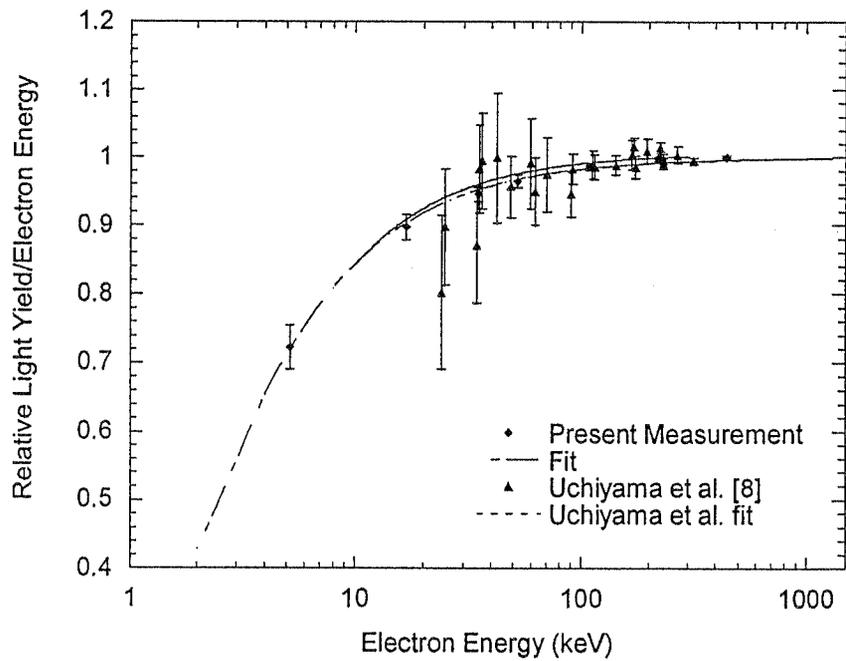


Fig. 2. GSO measured electron response. The response increases by nearly 28% as the electron energy increased from 5 keV to 445 keV.

YSO	$\text{Y}_2\text{SiO}_5:\text{Ce}$
GSO	$\text{Gd}_2\text{SiO}_5:\text{Ce}$
LSO	$\text{Lu}_2\text{SiO}_5:\text{Ce}$
LGSO	$\text{Lu}_{1-x}\text{Gd}_x\text{SiO}_5:\text{Ce}$

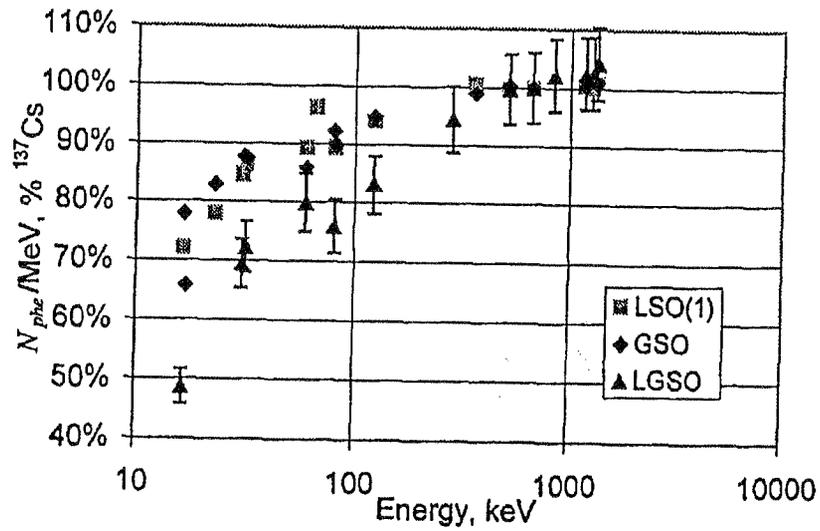


Figure 5. Nonproportionality of the light output for LSO, GSO and LGSO.

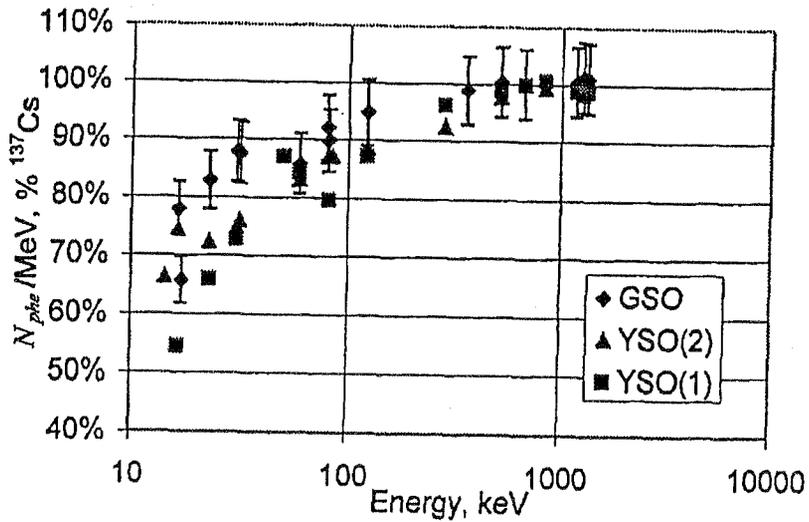


Figure 6. Nonproportionality of the light output for GSO and YSO (1) and (2).

## **Requirements for new scintillation detectors**

- high light yield
  - good energy resolution
  - fast
  - high stopping power
  - good mechanical and optical properties
- 

## **Ways to improve the scintillators**

- same crystal but with new dopants
- cooled scintillators
- new materials

	$\tau$ [ns]	$\lambda_{\max}$ [nm]	light yield [photons/MeV]
CsI pure	16	315	2 000
CsI(Tl)	800/6000	540	60 000
CsI(Na)	630	420	37 000
CsI(CO <sub>3</sub> )	2000	405	26 000

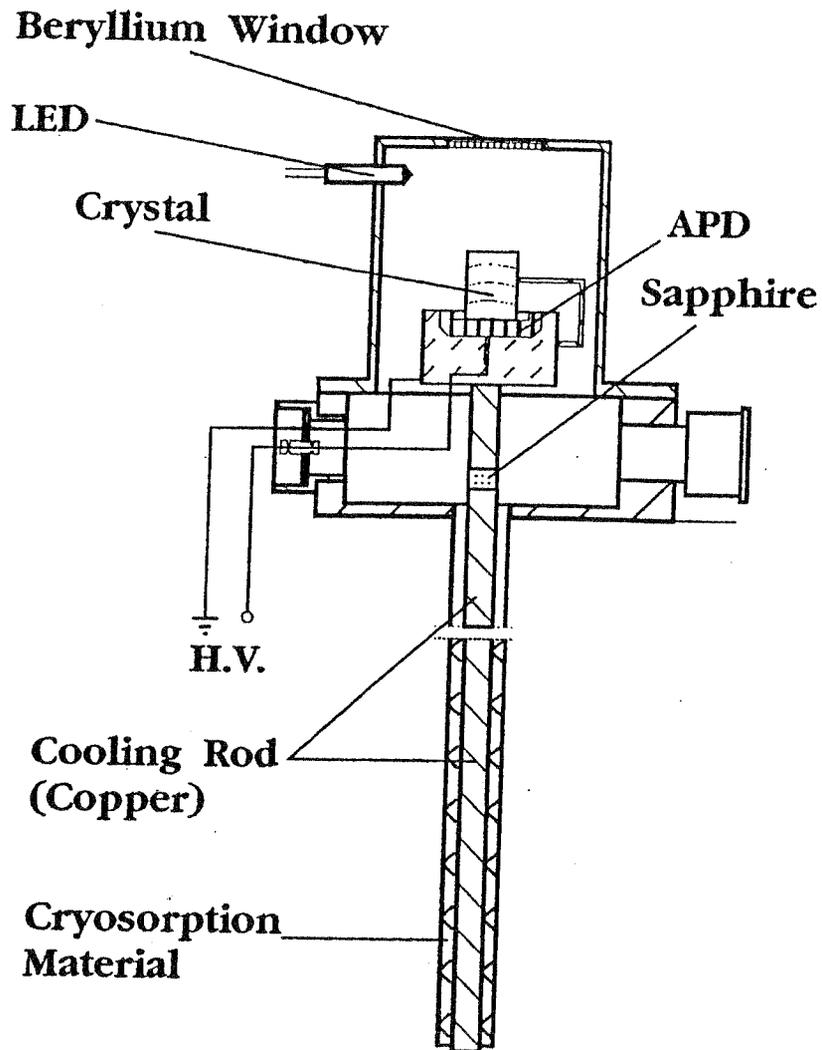
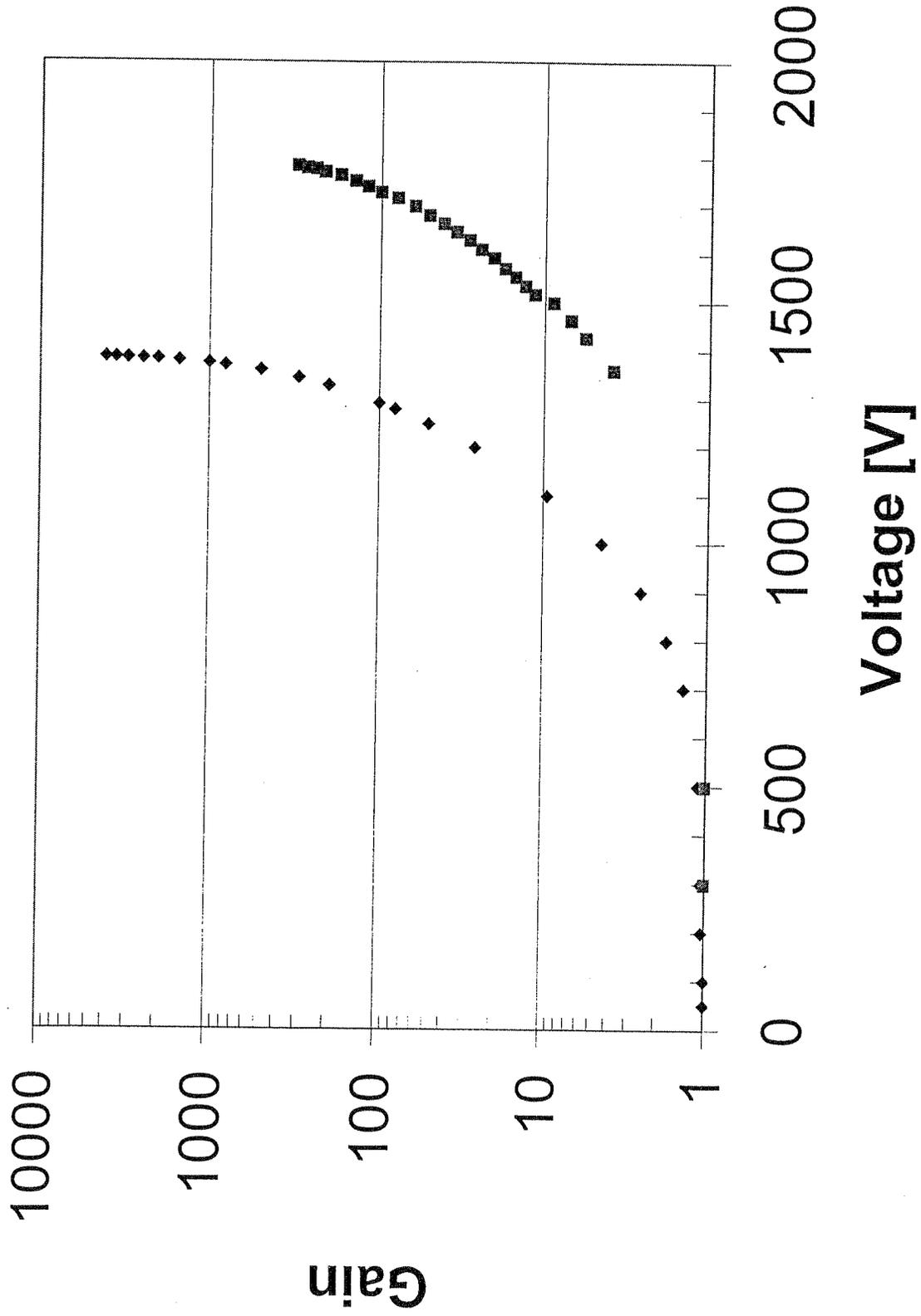
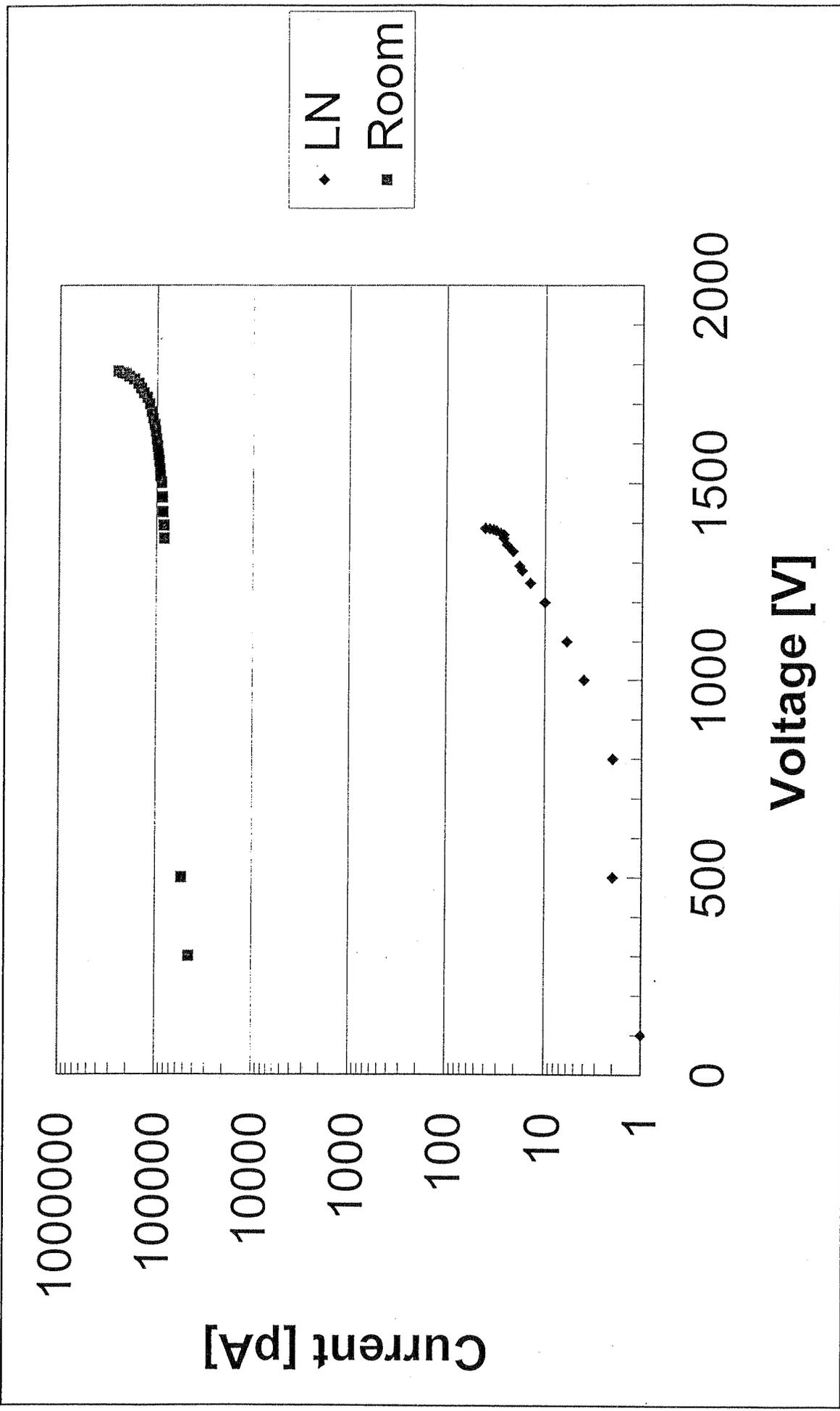


Fig. 1. Schematic drawing of the cryostat with the NaI crystal coupled to the LAAPD.



◆ LN  
■ Room



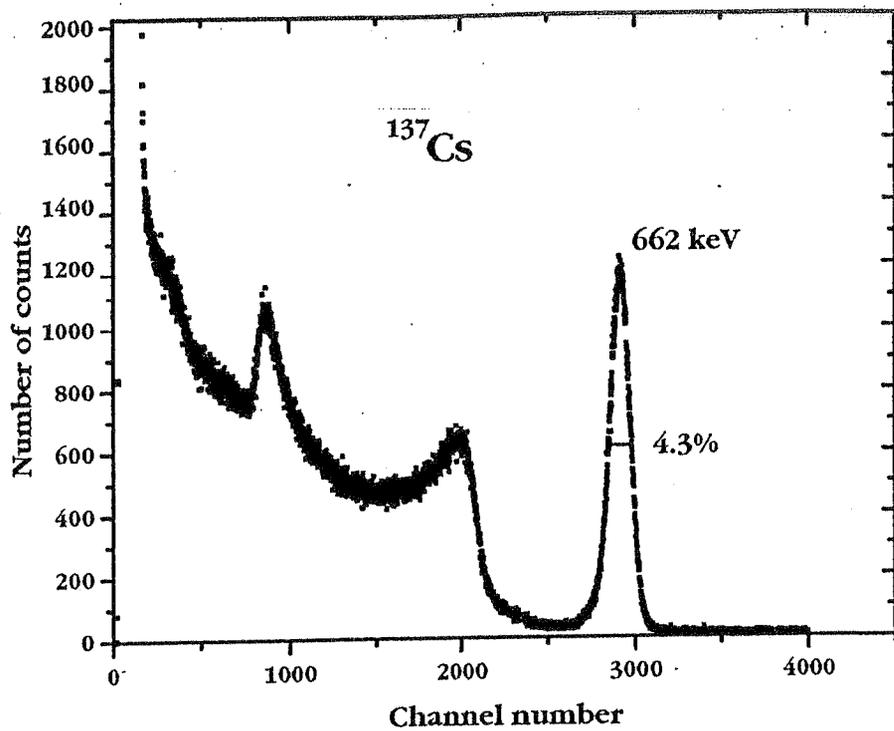


Fig. 6. Energy spectrum of  $\gamma$ -rays from a  $^{137}\text{Cs}$  source, as measured with the NaI (B) at  $\text{LN}_2$  temperature.

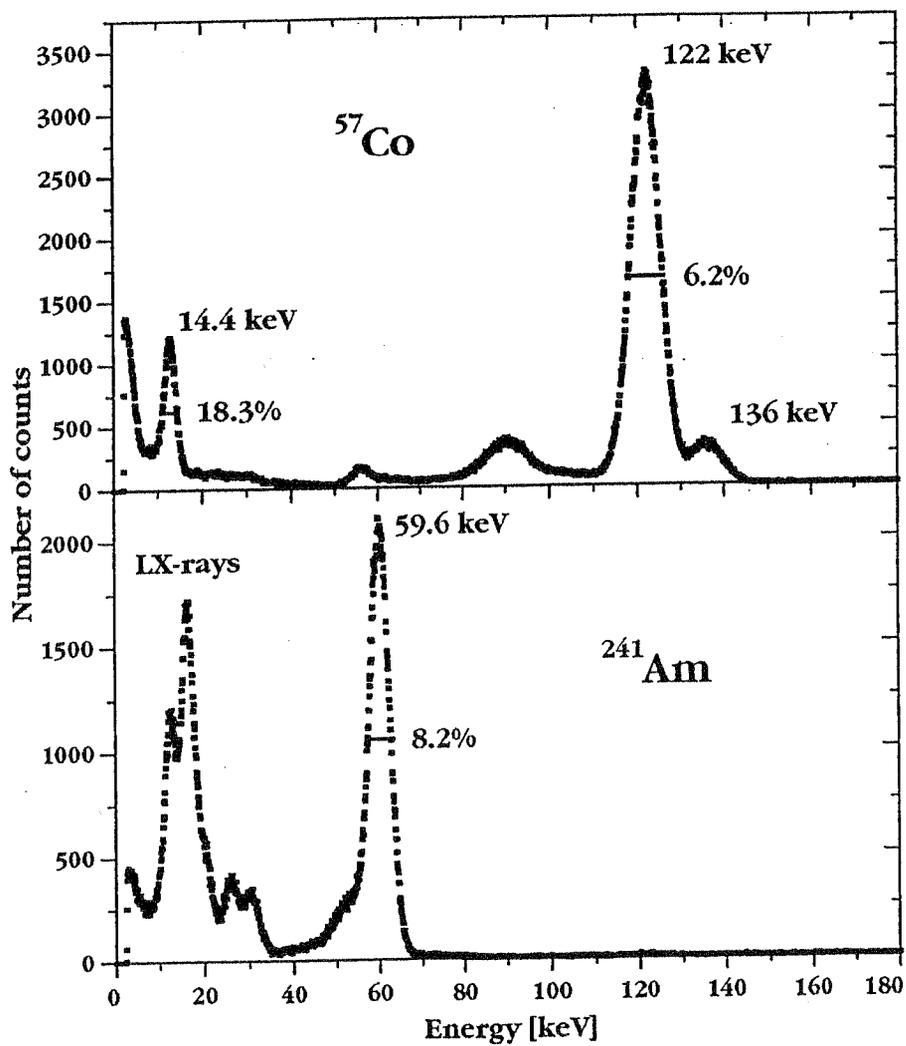


Fig. 7. Energy spectra of  $\gamma$ -rays from  $^{241}\text{Am}$  and  $^{57}\text{Co}$  sources measured with the NaI (B) at  $\text{LN}_2$  temperature.

## Pure NaI

crystal	<i>room temp.</i>		<i>LN<sub>2</sub> temp.</i>	
	energy resolution (%)	light yield (ph/MeV)	energy resolution (%)	light yield (ph/MeV)
A	16.2±1	4000±400	3.8±0.1	69000±7000
B	16.5±1	3800±400	4.3±0.2	84000±9000
C	14.9±1	2500±250	6.2±0.2	62000±7000

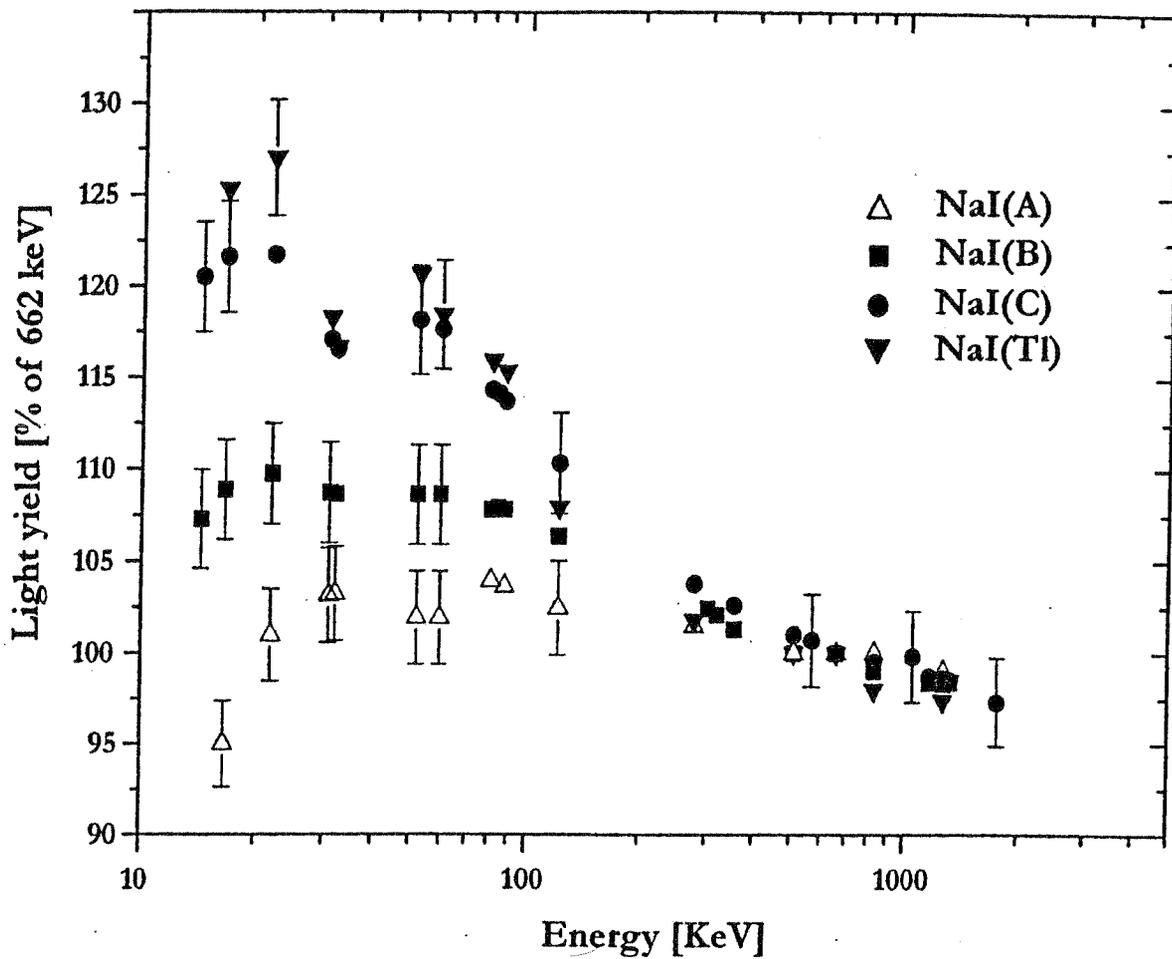


Fig. 8. The non-proportionality curves of studied NaI crystals in comparison to the curve measured for NaI(Tl) coupled to the XP2020Q photomultiplier. Error bars are shown for some selected points.

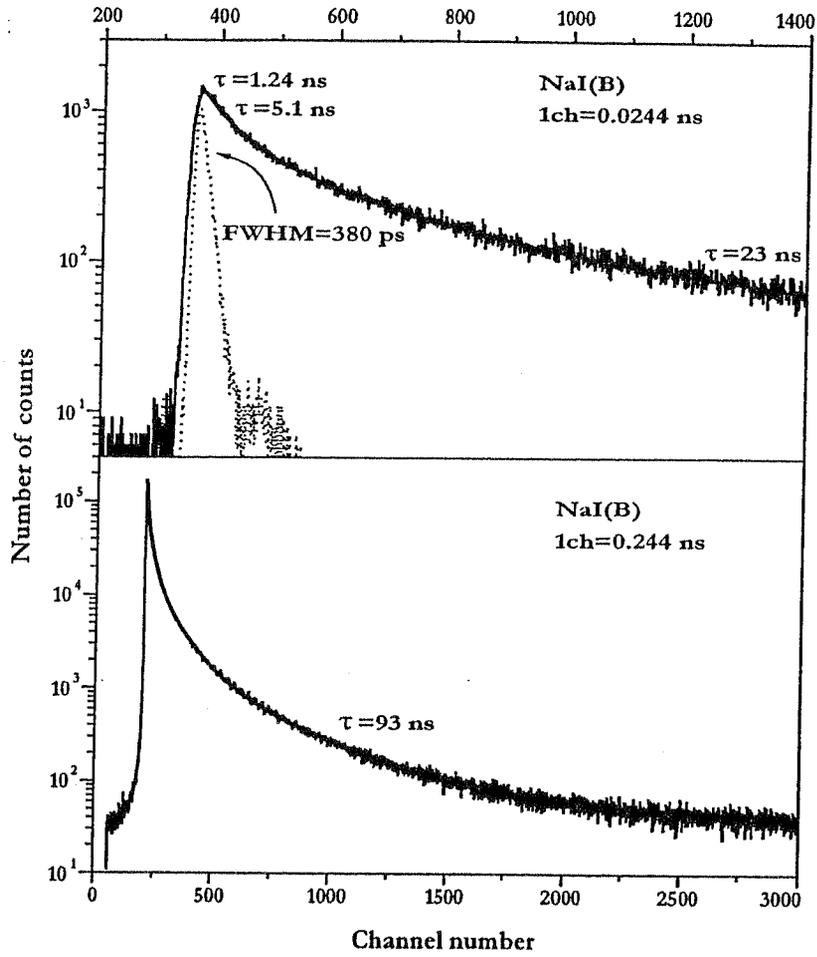


Fig. 4. Light pulse shape from a pure NaI measured at room temperature is shown in two different time intervals. The prompt spectrum of the timing system is shown in the upper plot.

TABLE II  
LIGHT PULSE COMPONENTS OF PURE NaI AT ROOM  
TEMPERATURE

Crystal	Decay time constants/ intensities			
A	$0.86 \pm 0.04$ ns /7.3 $\pm$ 0.4%	$4.5 \pm 0.1$ ns /27.4 $\pm$ 0.9%	$22.3 \pm 0.3$ ns /51.5 $\pm$ 1.7%	$120 \pm 3$ ns /13.8 $\pm$ 0.8%
B	$1.24 \pm 0.05$ ns /11.1 $\pm$ 0.7%	$5.1 \pm 0.2$ ns /27.6 $\pm$ 2.4%	$23 \pm 1$ ns /44.4 $\pm$ 4.1%	$93 \pm 1$ ns /16.9 $\pm$ 0.3%
C	$1.02 \pm 0.03$ ns /15.2 $\pm$ 0.5%	$4.2 \pm 0.1$ ns /30.1 $\pm$ 0.9%	$18.6 \pm 0.3$ ns /46.9 $\pm$ 1.4%	$138 \pm 8$ ns /7.8 $\pm$ 0.6%

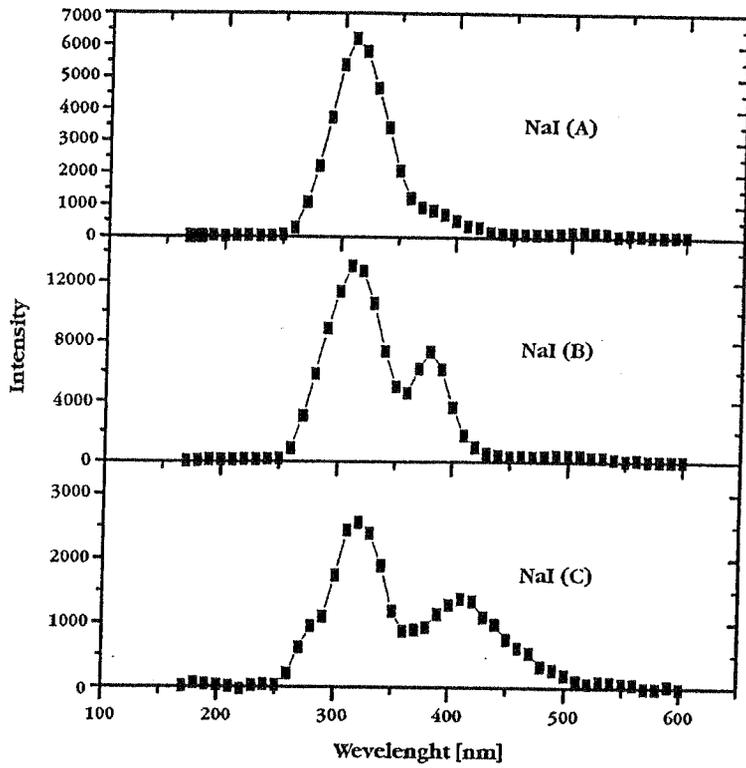


Fig. 2. Emission spectra of tested samples of pure NaI crystal.

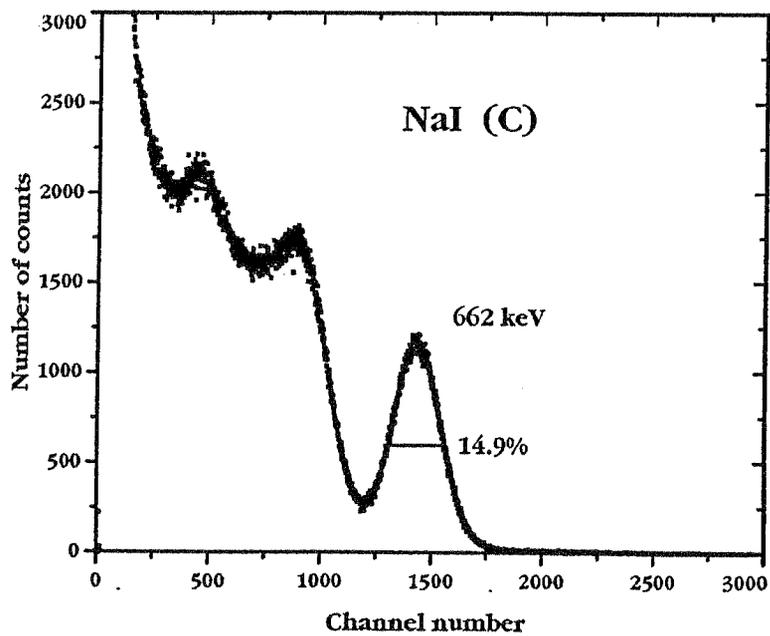


Fig. 3. Energy spectrum of a  $^{137}\text{Cs}$  source, as measured with NaI (C).

**Room temp.**

	$\tau_d$ [ns]	$\lambda_{\max}$ [nm]	light yield [ph/MeV]
CsI(Tl)	800/6000	540	60 000
CsI pure	16	315	2 000

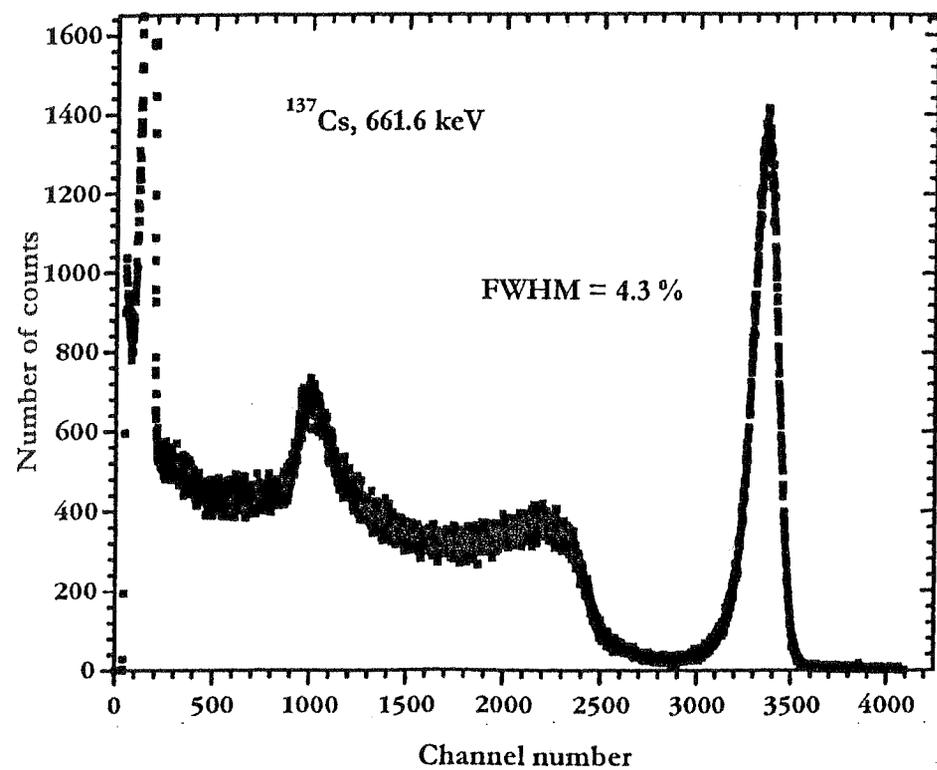


Fig. 2. Energy spectrum of 662 keV  $\gamma$ -rays from a  $^{137}\text{Cs}$  source measured with CsI (B).

Table 1

Number of e-h pairs, light output and energy resolution of CsI crystals measured at 662 keV  $\gamma$ -rays

Crystal	$N_{e-h}/\text{MeV}$	$\Delta E/E$ [%]	Light output [ph/MeV]
CsI - A	$37600 \pm 800$	$5.0 \pm 0.2\%$	$108000 \pm 10000$
CsI - B	$43000 \pm 900$	$4.3 \pm 0.1\%$	$124000 \pm 12000$

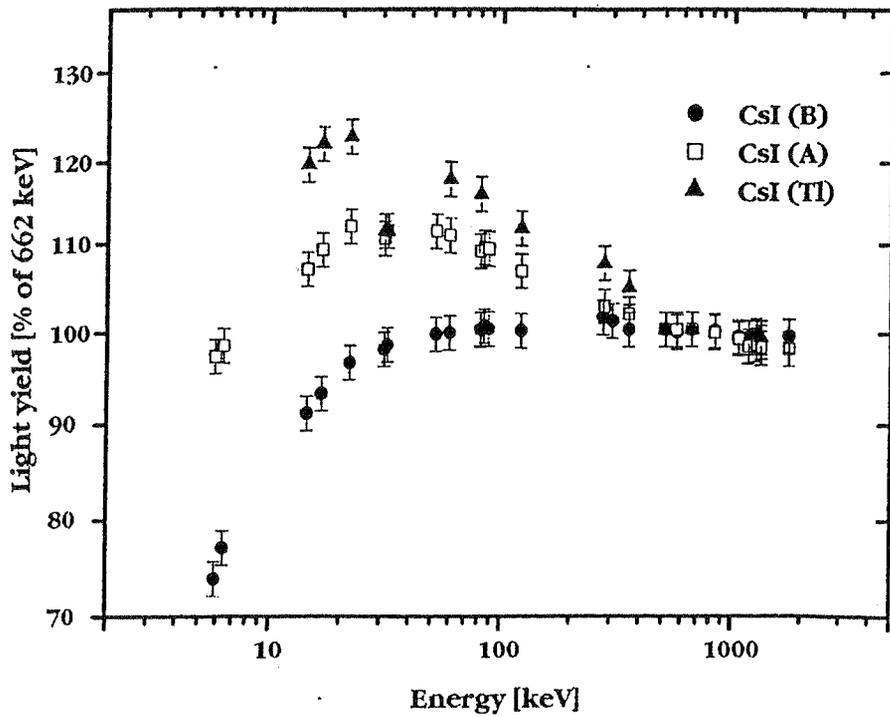


Fig. 3. The non-proportionality curves of studied CsI crystals at LN<sub>2</sub> temperature in comparison to the curve measured for CsI(Tl) coupled to the XP2020Q photomultiplier at room temperature.

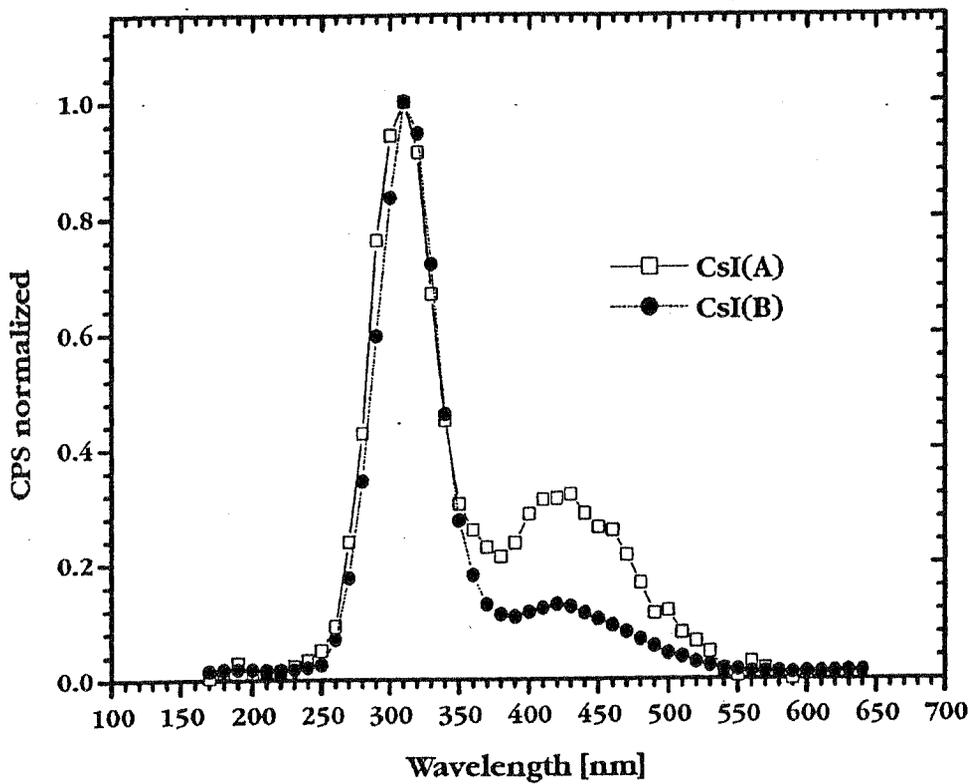


Fig. 8. Emission spectra of tested pure CsI crystals, as measured at room temperature.

## BGO ( $\text{Bi}_4\text{Ge}_3\text{O}_{12}$ )

Crystals diam. 9x4 mm

crystal	<i>room temp.</i>		<i>LN<sub>2</sub> temp.</i>	
	energy resolution (%)	light yield (ph/MeV)	energy resolution (%)	light yield (ph/MeV)
BGO 1	10.0±0.3	6900±140	6.5±0.2	29000±2000
BGO 2			6.3±0.2	29500±2000

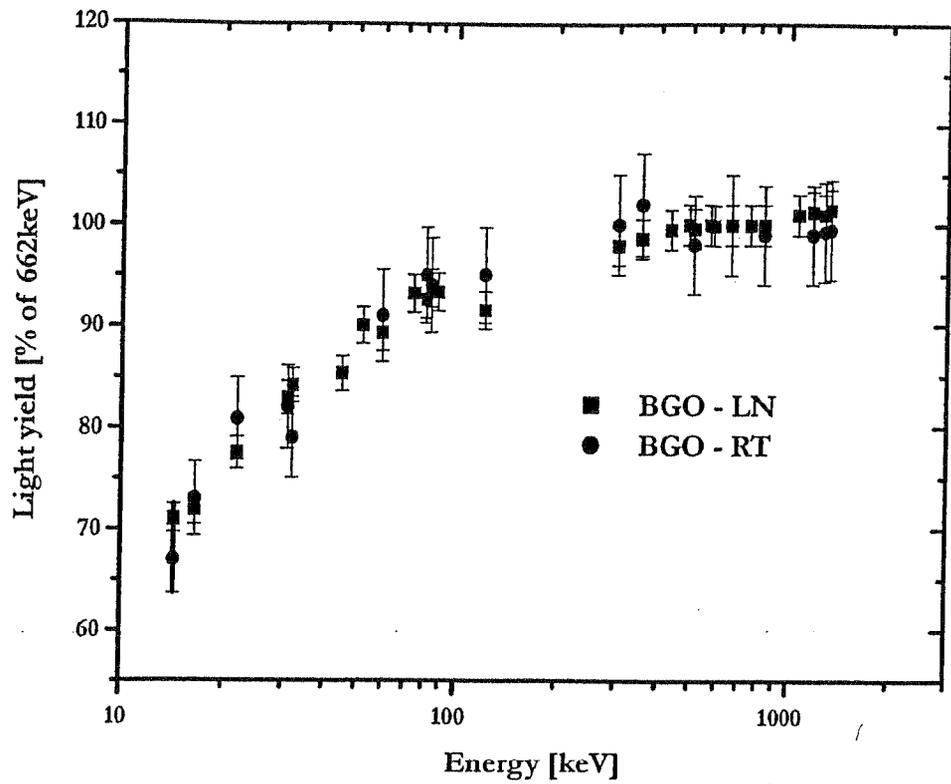


Fig. 6. The non-proportionality characteristics of BGO crystals determined at both room and LN<sub>2</sub> temperatures.

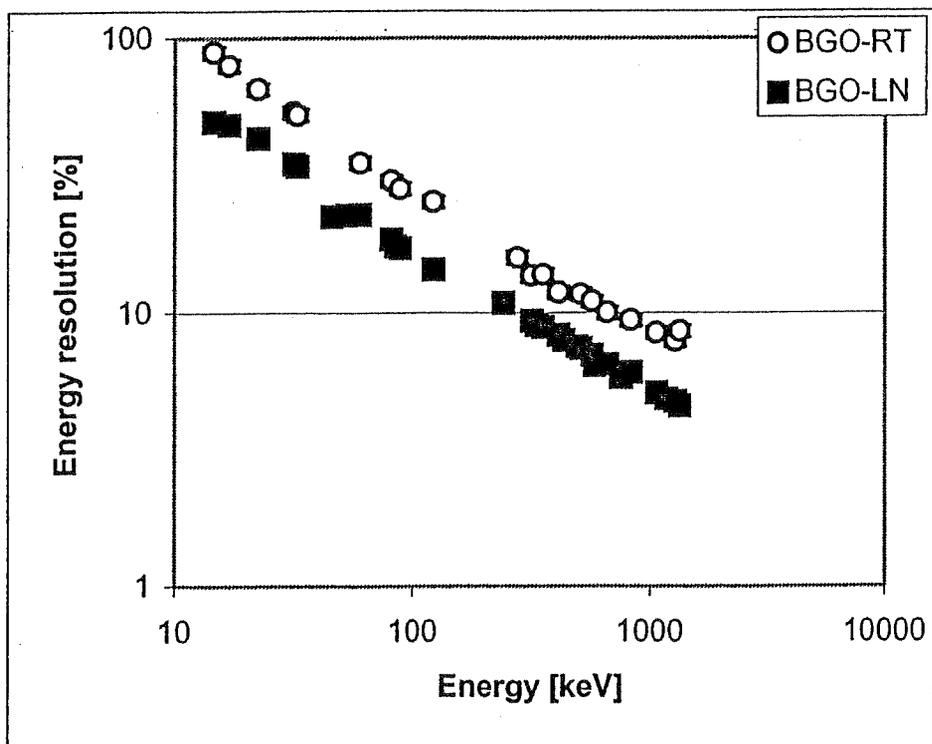
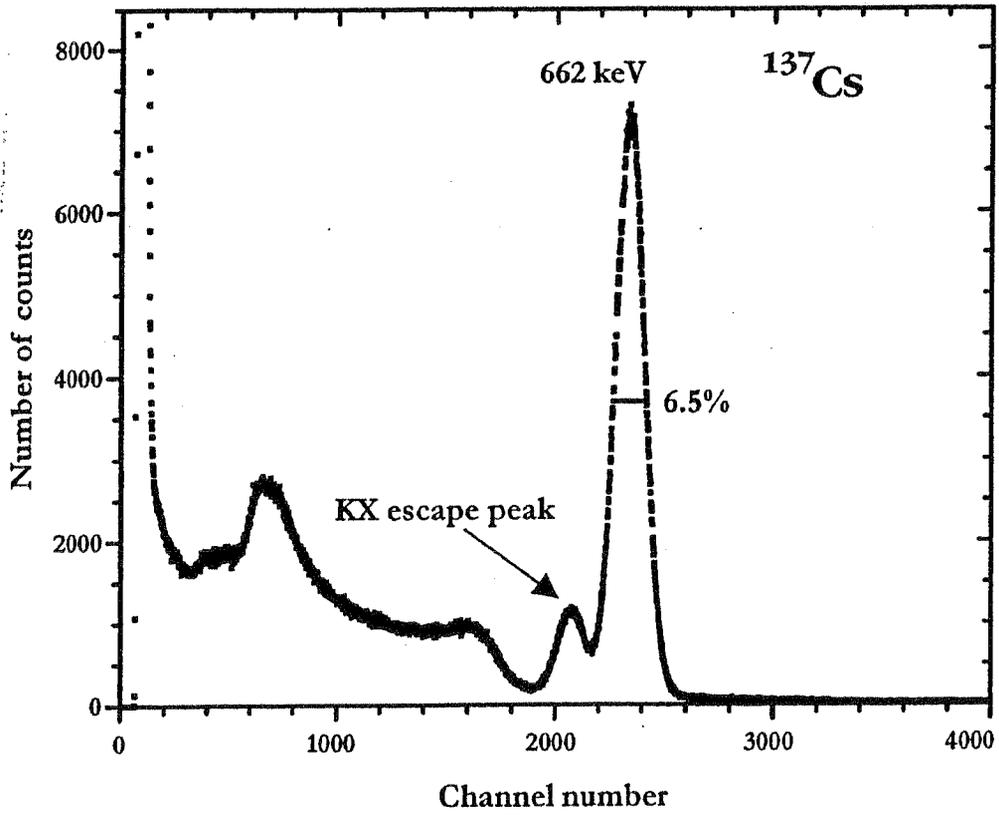
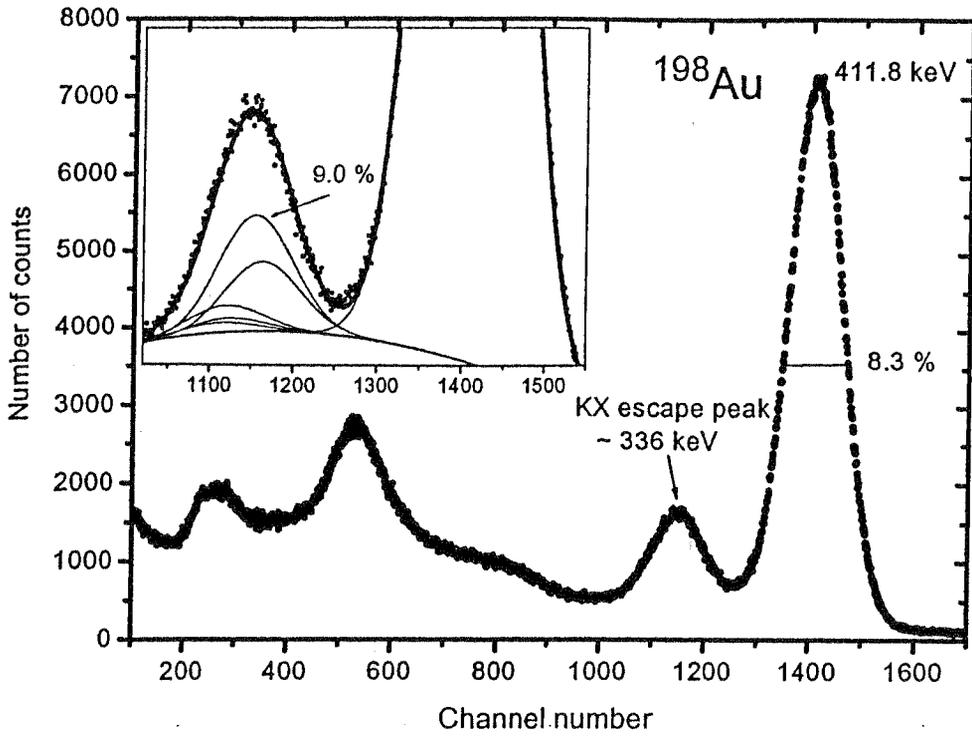


Fig. 7. Energy resolution measured with BGO at room and LN<sub>2</sub> temperature versus energy of gamma rays.



**LaBr<sub>3</sub>:Ce**  
(0.5 % Ce)

density: 5.29 g/cm<sup>3</sup>

hygroscopic

$\lambda_{\text{max}}=358$  nm

$\tau=35$  ns

light output: 61 000 ph/MeV

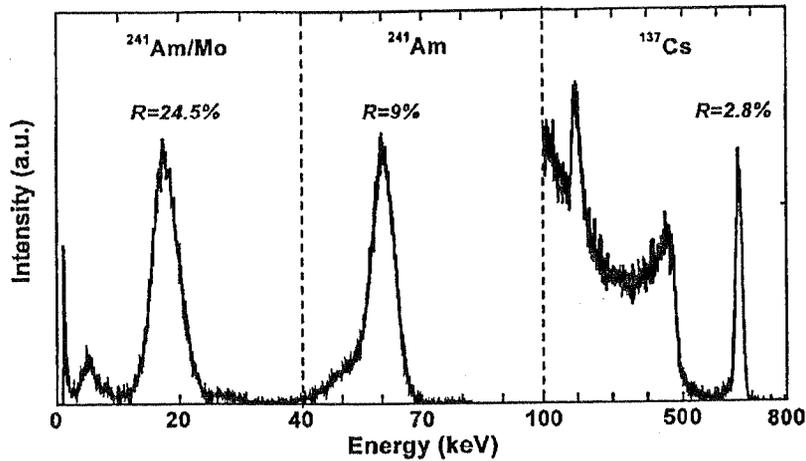


FIG. 1. Energy resolution obtained with a single crystal of LaBr<sub>3</sub>:0.5% Ce<sup>3+</sup> for x/γ rays from <sup>137</sup>Cs, <sup>241</sup>Am, and <sup>241</sup>Am/Mo source.

# LaCl<sub>3</sub>:Ce

density: 3.86 g/cm<sup>3</sup>

hygroscopic

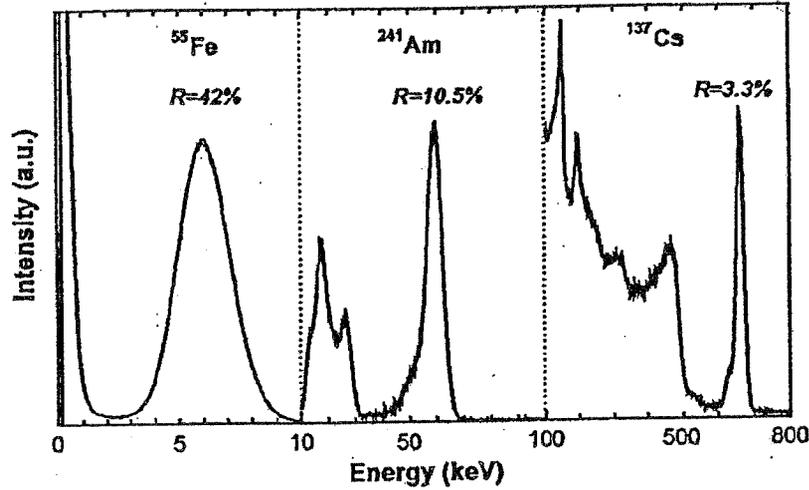


FIG. 1. Energy resolutions obtained with a single crystal of LaCl<sub>3</sub>:Ce 10% for x/γ rays from (a) <sup>55</sup>Fe, (b) <sup>241</sup>Am, and (c) <sup>137</sup>Cs sources.

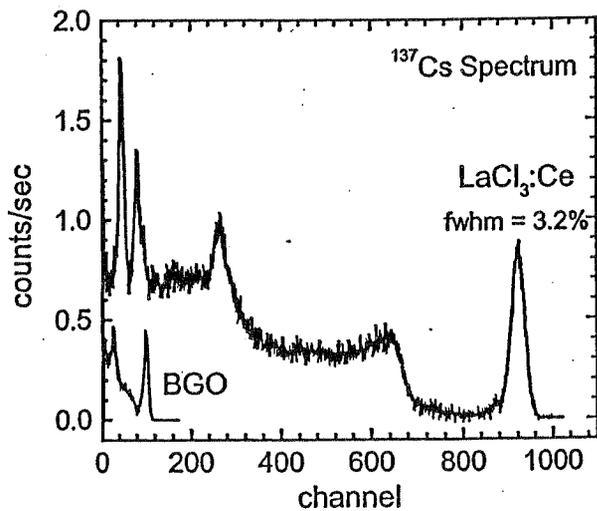


Fig. 1. <sup>137</sup>Cs spectra recorded with LaCl<sub>3</sub>:10% Ce and BGO crystals coupled to PMT under identical operating conditions. Light output of LaCl<sub>3</sub>:10% Ce was estimated to be ~50,000 photons/MeV from the calibration provided by BGO. The LaCl<sub>3</sub> spectrum shows resolution of 3.2% (FWHM) at room temperature.

LaCl<sub>3</sub>:Ce

% Ce	L.Y [photons/MeV]	$\tau$ [ns]	$\lambda$ [nm]
0.1	50 500	20 (15%), 213 (85%)	350, 430
1	50 500	20 (33%), 213 (67%)	350, 430
10	50 000	20 (70%), 213 (30%)	350, 430
20	38 000	25 (76%), 63 (13%), 213 (11%)	350, 430

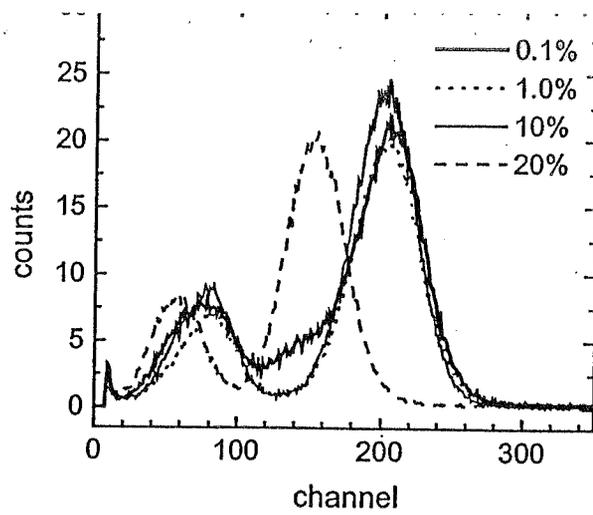


Fig. 2.  $^{241}\text{Am}$  spectra (60 keV photons) recorded with  $\text{LaCl}_3:\text{Ce}$  crystals of different Ce concentrations. From the peak position light output was estimated for each crystal, see Table 1.

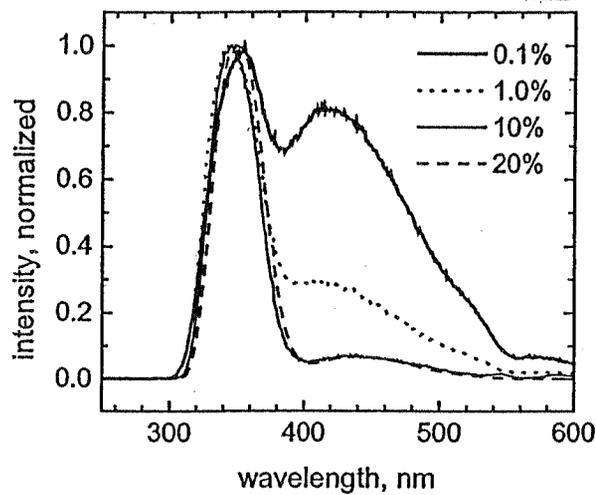


Fig. 3. Optical emission spectra for  $\text{LaCl}_3$  samples with different Ce concentrations. Spectra were normalized with respect to 350 nm peak intensity.

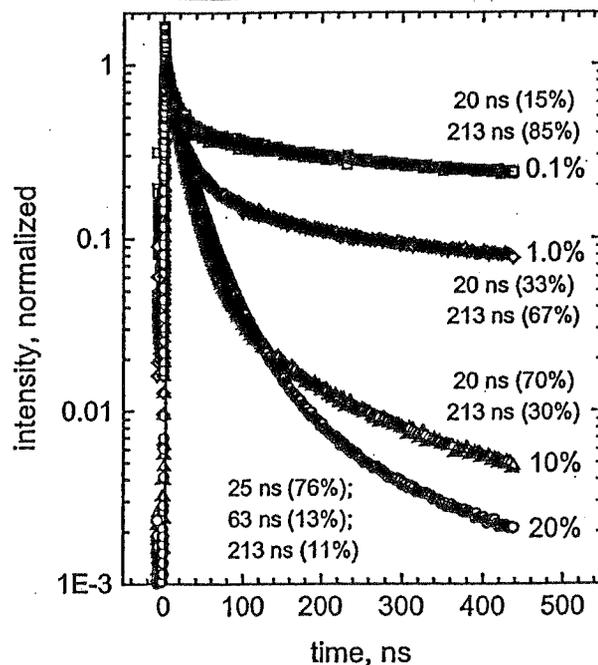


Fig. 4. Time profiles (points) measured for  $\text{LaCl}_3$  crystals with 0.1%, 1.0%, 10%, and 20% Ce concentrations, along with multi-exponential fits (lines). As seen in plots, the principal decay constant is fast ( $\leq 25$  ns) for samples with high (10–20%) Ce concentration.

**LYSO**  
**LuYSiO<sub>5</sub>:Ce**

density: 7.1 g/cm<sup>3</sup>

$\lambda_{\text{max}}=430$  nm

$\tau=48$  ns

light output: 32 000 photons/MeV

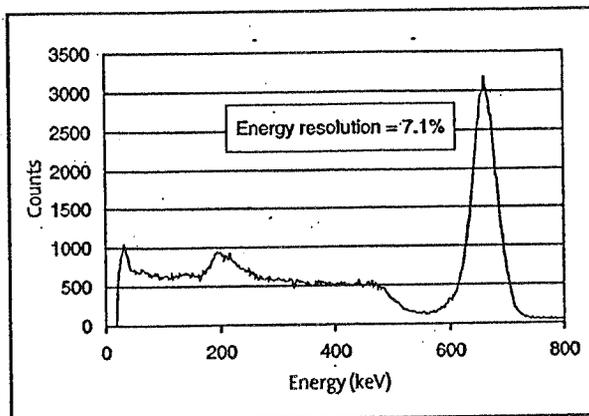


Figure 1. LYSO Response to 662 keV Photons

	L.Y [ph/MeV]	energy resolution [%]	time resolution [ps]
LaBr <sub>3</sub> :Ce	61 000	2.8	385
LaCl <sub>3</sub> :Ce	50 000	3.2	224
LYSO:Ce	32 000	7.1	-