



LSO/LYSO Crystal Development

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Why Crystal Calorimeter



- Enhance physics discovery potential since photons and electrons are fundamental particles for the standard model physics and new physics.
- Performance of a crystal calorimeter is well understood:
 - The best possible energy resolution, good position and photon angular resolution;
 - Good e/photon identification and reconstruction efficiency;
 - Good missing energy resolutions;
 - Good jet energy resolution.



Physics with Crystal Calorimeters

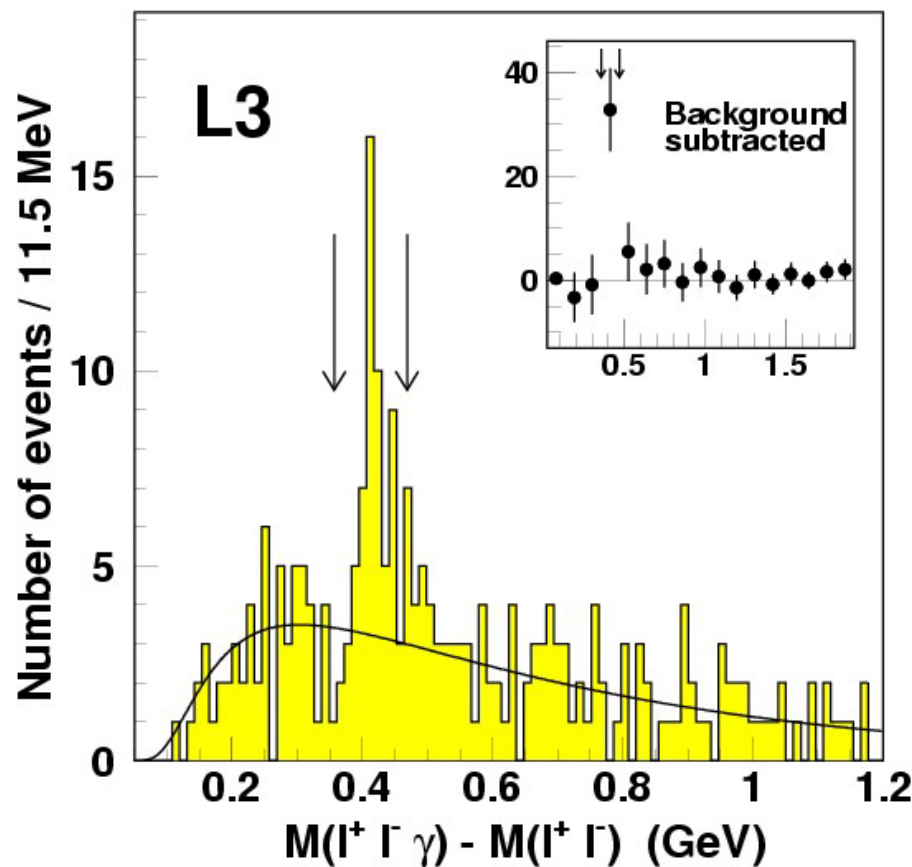
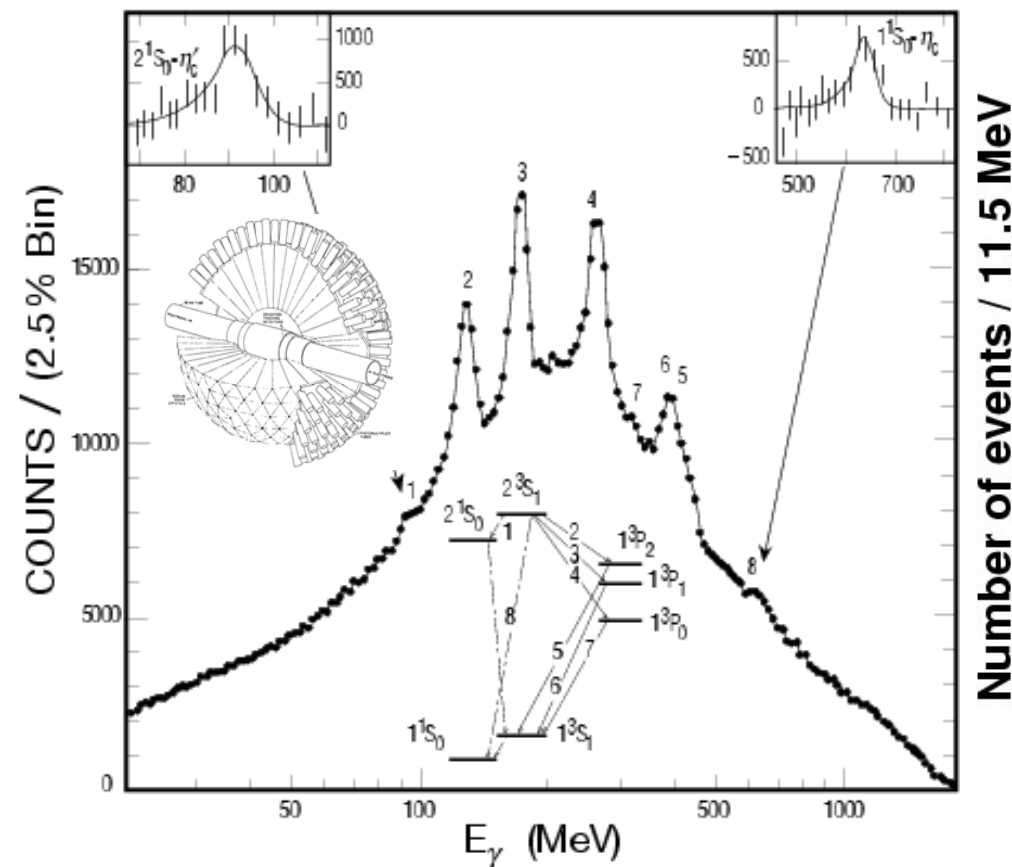
Charmonium system observed
by CB through Inclusive photons

CB NaI(Tl)

Charmed Meson in Z Decay

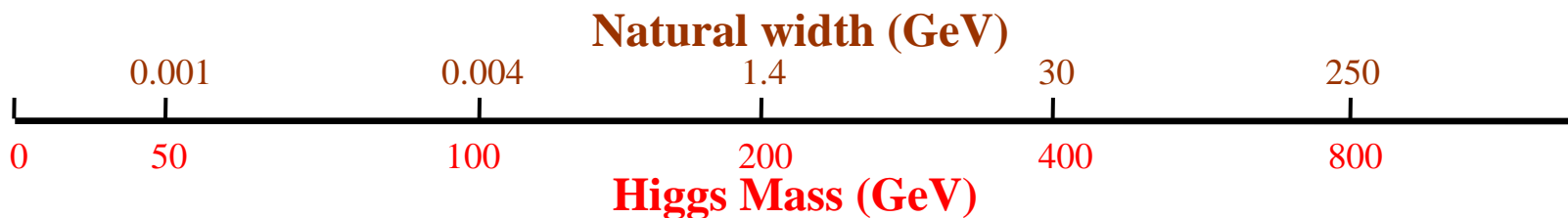
$$\chi_{c1} \rightarrow J/\psi \gamma$$

L3 BGO





H $\rightarrow\gamma\gamma$ Search Needs Precision ECAL



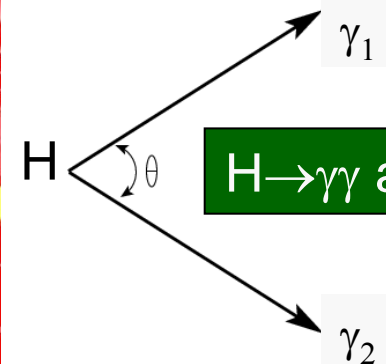
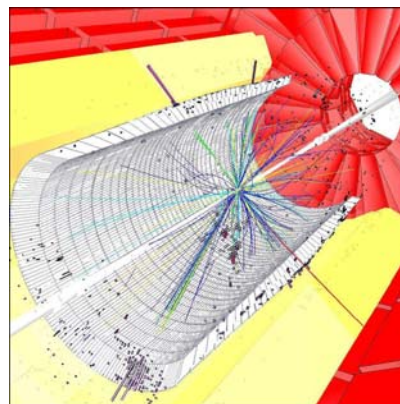
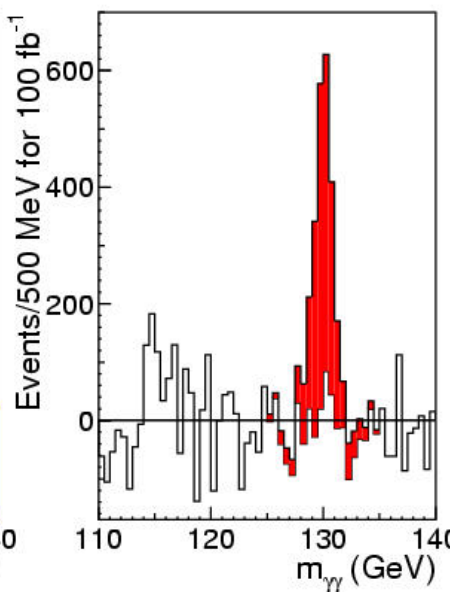
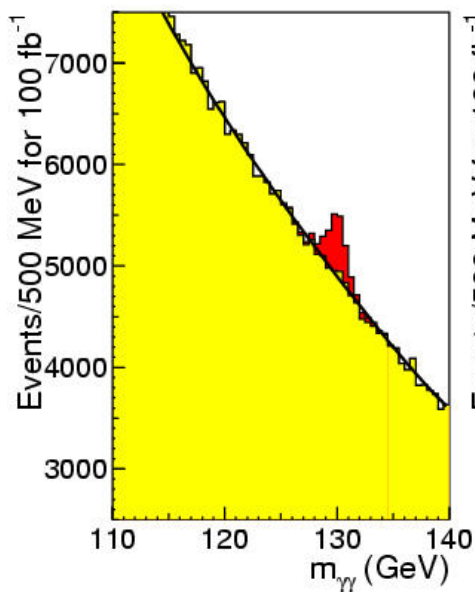
H \rightarrow ZZ* \rightarrow 4 leptons

CMS PWO

Narrow width and large background

H \rightarrow ZZ \rightarrow 4 leptons

H \rightarrow WW or ZZjj

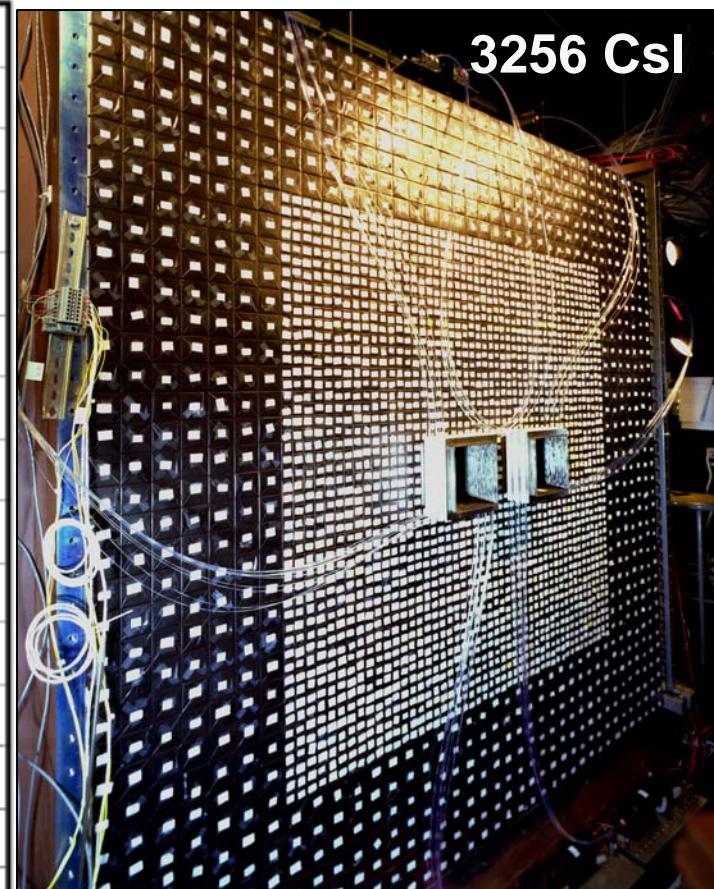
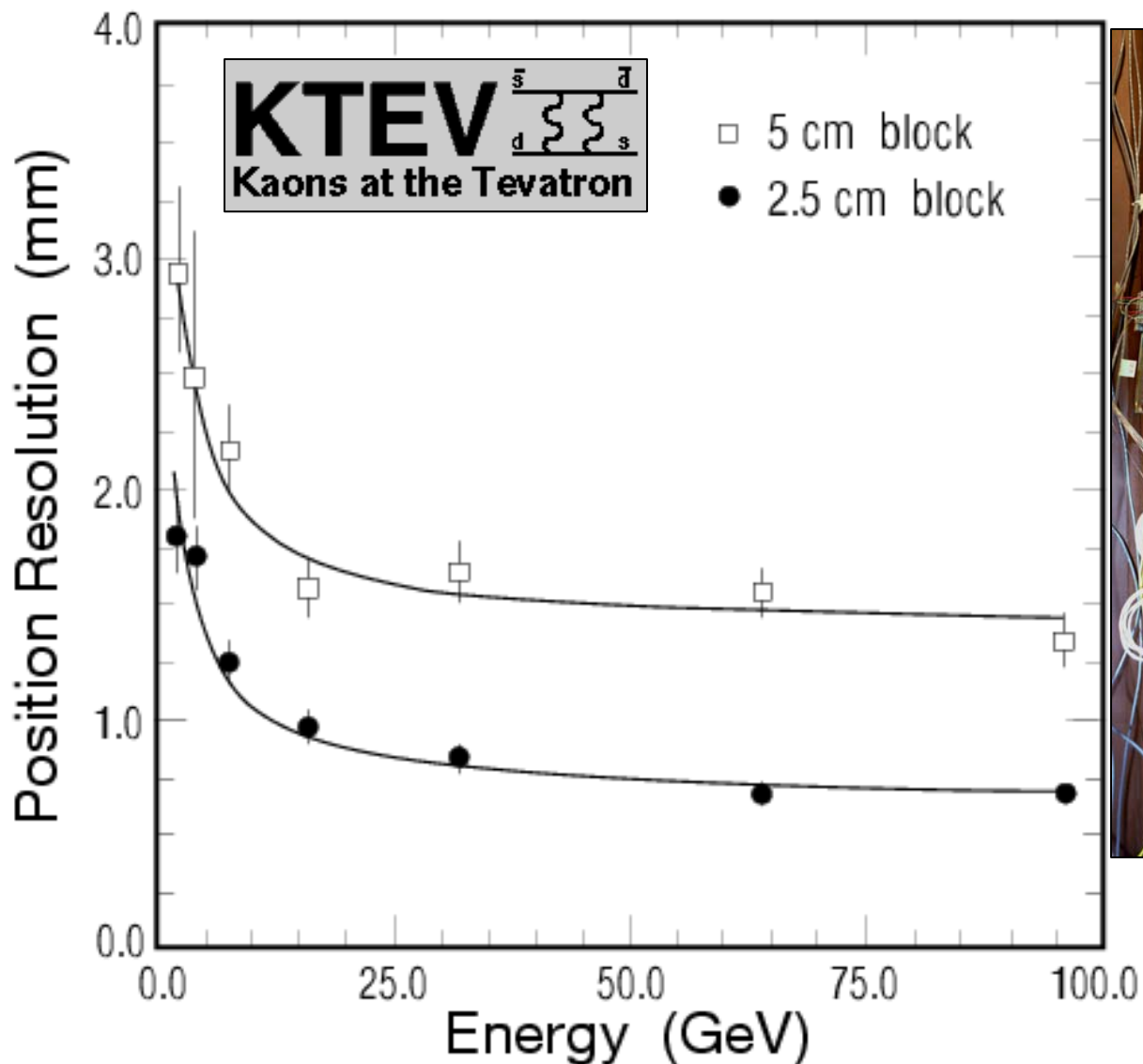


H $\rightarrow\gamma\gamma$ at LHC

$$\sigma m / m = 0.5 [\sigma E_1 / E_1 \oplus \sigma E_2 / E_2 \oplus \sigma \theta / \tan(\theta/2)],$$

where $\sigma E / E = a / \sqrt{E} \oplus b \oplus c/E$ and E in GeV

KTeV CsI Position Resolution

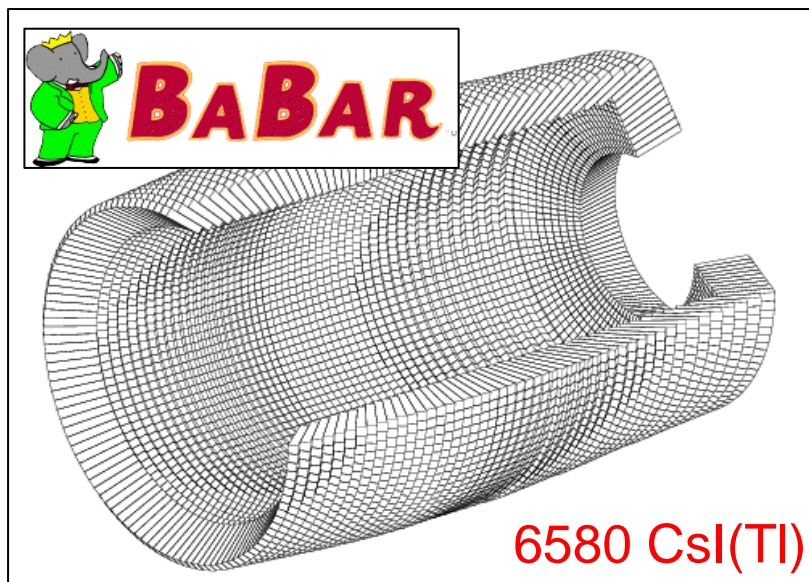


$$\sigma_x = 3 / \sqrt{E} \oplus 0.3 \text{ mm for L3 BGO \& CMS PWO}$$

BaBar CsI(Tl) Energy Resolution



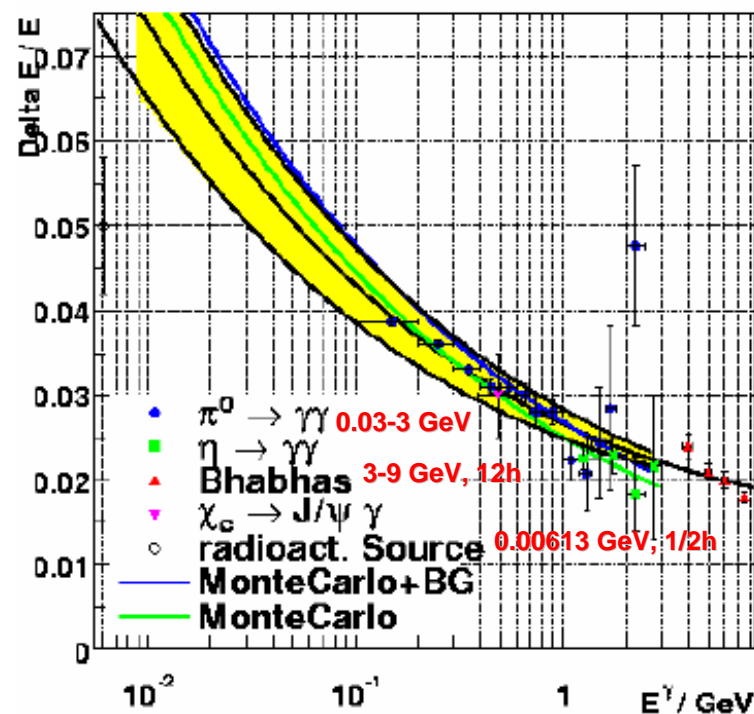
BaBar CsI(Tl)



Good light yield of CsI(Tl)
provides excellent energy
resolution at low energies

Energy resolution

M. Kocian, SLAC, CALOR2002



$$\frac{\sigma_E}{E} = \frac{\sigma_1}{\sqrt{E}} \oplus \sigma_2$$

$$\sigma_1 = (2.30 \pm 0.03 \pm 0.3)\%$$

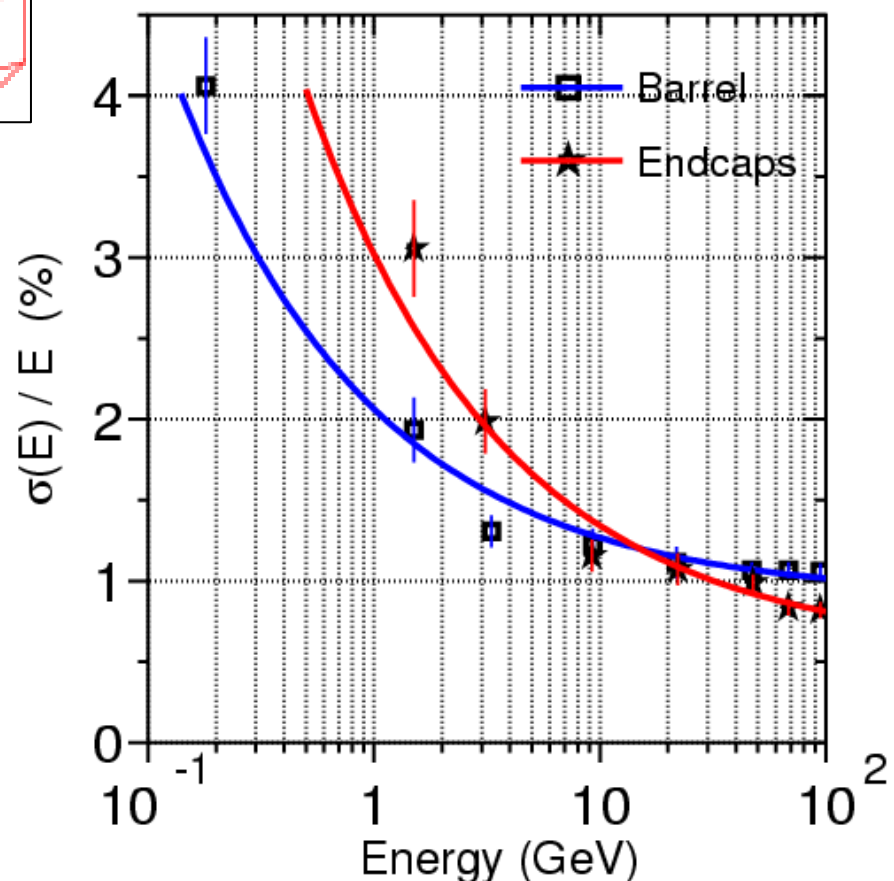
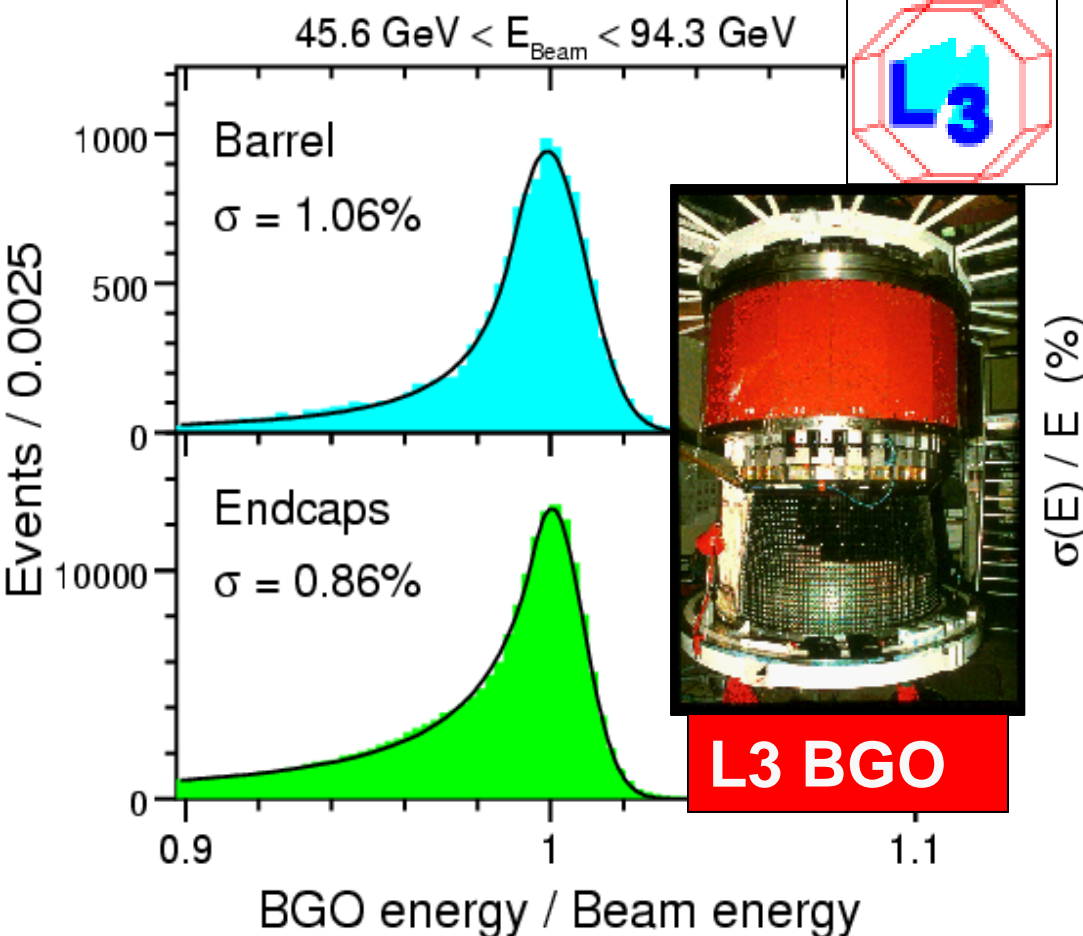
$$\sigma_2 = (1.35 \pm 0.08 \pm 0.2)\%$$



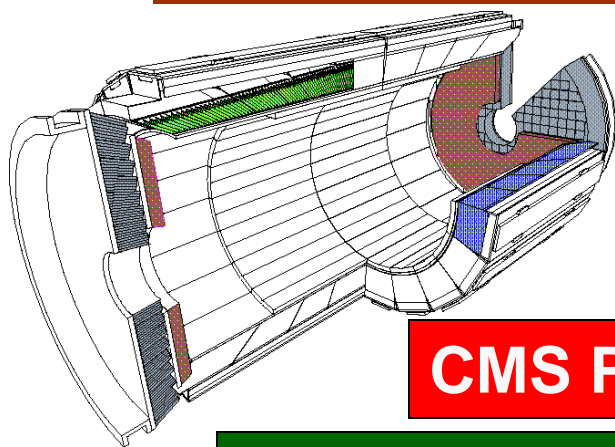
L3 BGO Energy Resolution



Contribution	"Radiative"+Intrinsic	Temperature	Calibration	Overall
Barrel	0.8%	0.5%	0.5%	1.07%
Endcaps	0.6%	0.5%	0.4%	0.88%



CMS PWO Energy Resolution



CMS PWO

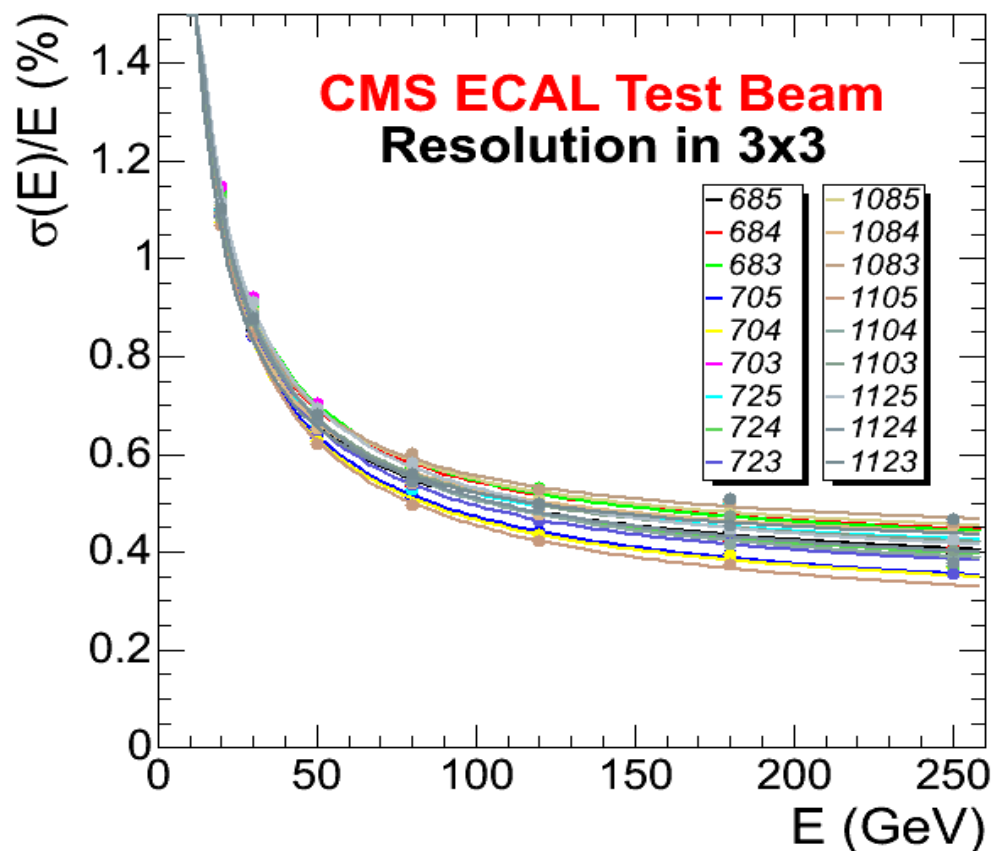
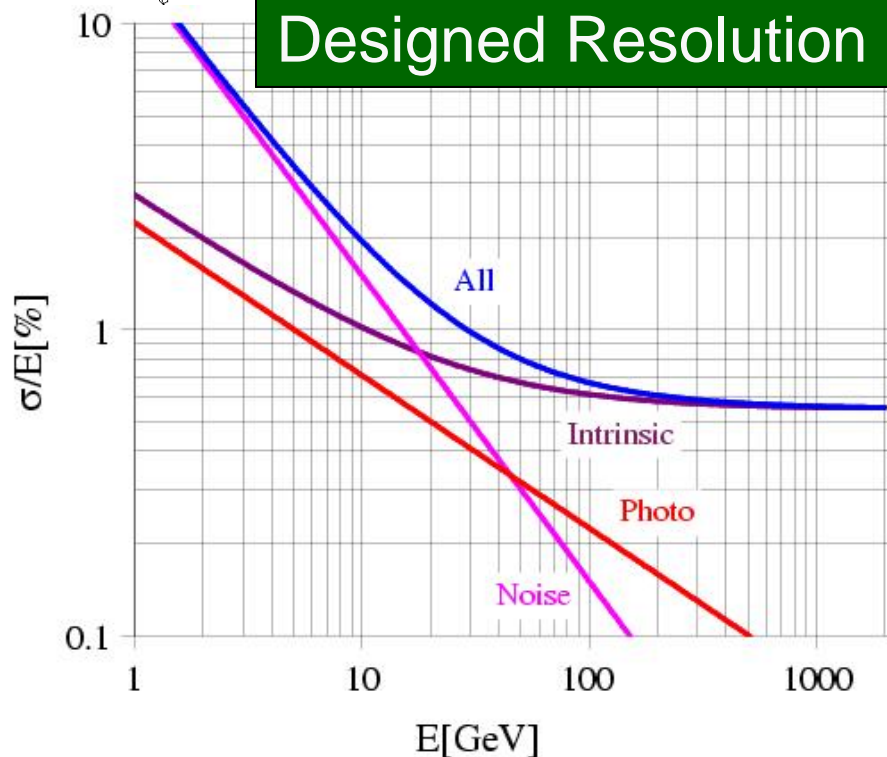


Measured Resolution

$$\sigma(E)/E < 1\% \text{ if } E > 25 \text{ GeV}$$

$$\sigma(E)/E \sim 0.5\% \text{ at } 120 \text{ GeV}$$

Designed Resolution





Bright Fast Dense LSO Crystal

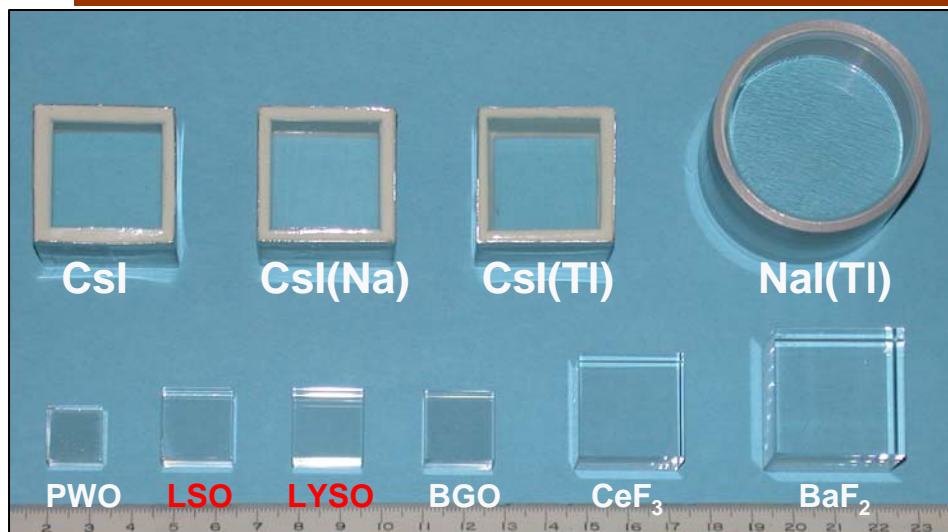


Crystal	Nal(Tl)	Csl(Tl)	Csl(Na)	Csl	CeF ₃	BaF ₂	BGO	PWO(Y)	LSO(Ce)
Density (g/cm ³)	3.67	4.51	4.51	4.51	6.16	4.89	7.13	8.3	7.40
Melting Point (°C)	651	621	621	621	1460	1280	1050	1123	2050
Radiation Length (cm)	2.59	1.86	1.86	1.86	1.65	2.03	1.12	0.89	1.14
Molière Radius (cm)	4.13	3.57	3.57	3.57	3.38	3.10	2.23	2.00	2.07
Interaction Length (cm)	42.9	39.3	39.3	39.3	23.17	30.7	22.8	20.7	20.9
Refractive Index ^a	1.85	1.79		1.95	1.62	1.50	2.15	2.20	1.82
Hygroscopicity	Yes	Slight	Slight	Slight	No	No	No	No	No
Luminescence ^b (nm) (at peak)	410	550	420	420 310	340 300	300 220	480	425 420	402
Decay Time ^b (ns)	245	1220	690	30 6	30	650 0.9	300	30 10	40
Light Yield ^{b,c} (%)	100	165	88	3.6 1.1	7.3	36 4.1	21	0.3 0.1	85
d(LY)/dT ^b (%/°C)	-0.2	0.4	0.4	-1.4	0	-1.9 0.1	-0.9	-2.5	-0.2

a. at peak of emission; b. up/low row: slow/fast component; c. QE of readout device taken out.



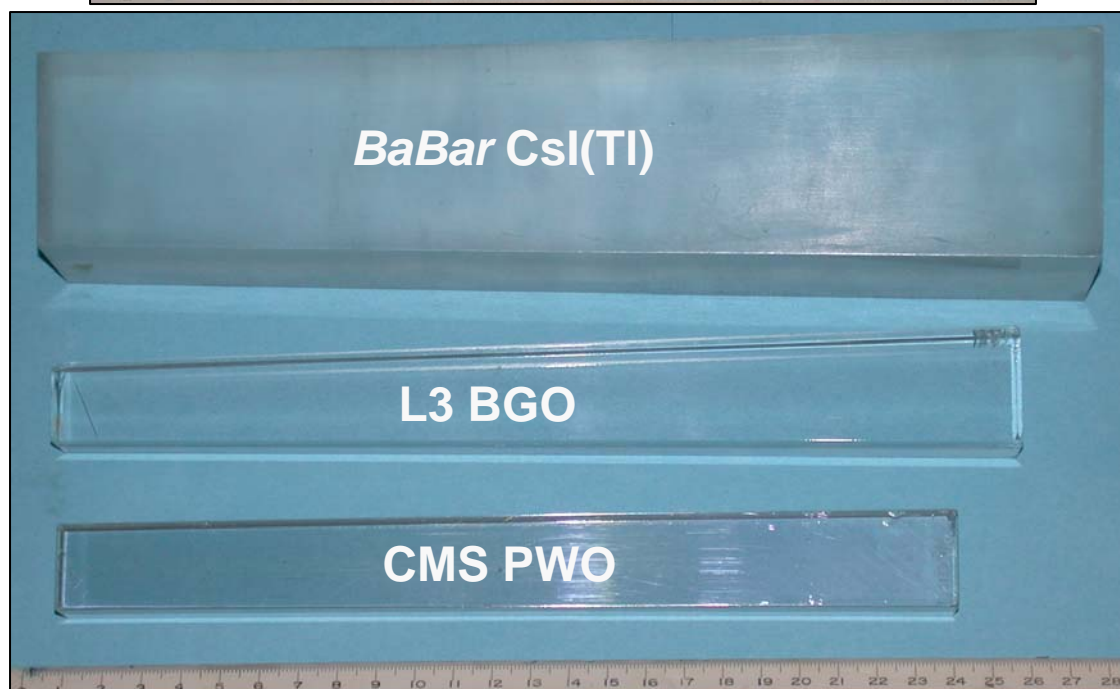
Crystal Density: Radiation Length



1.5 X_0 Cubic Samples:

Hygroscopic Halides

Non-hygroscopic



Full Size Crystals:

BaBar Csl(Tl): 16 X_0

L3 BGO: 22 X_0

CMS PWO(Y): 25 X_0

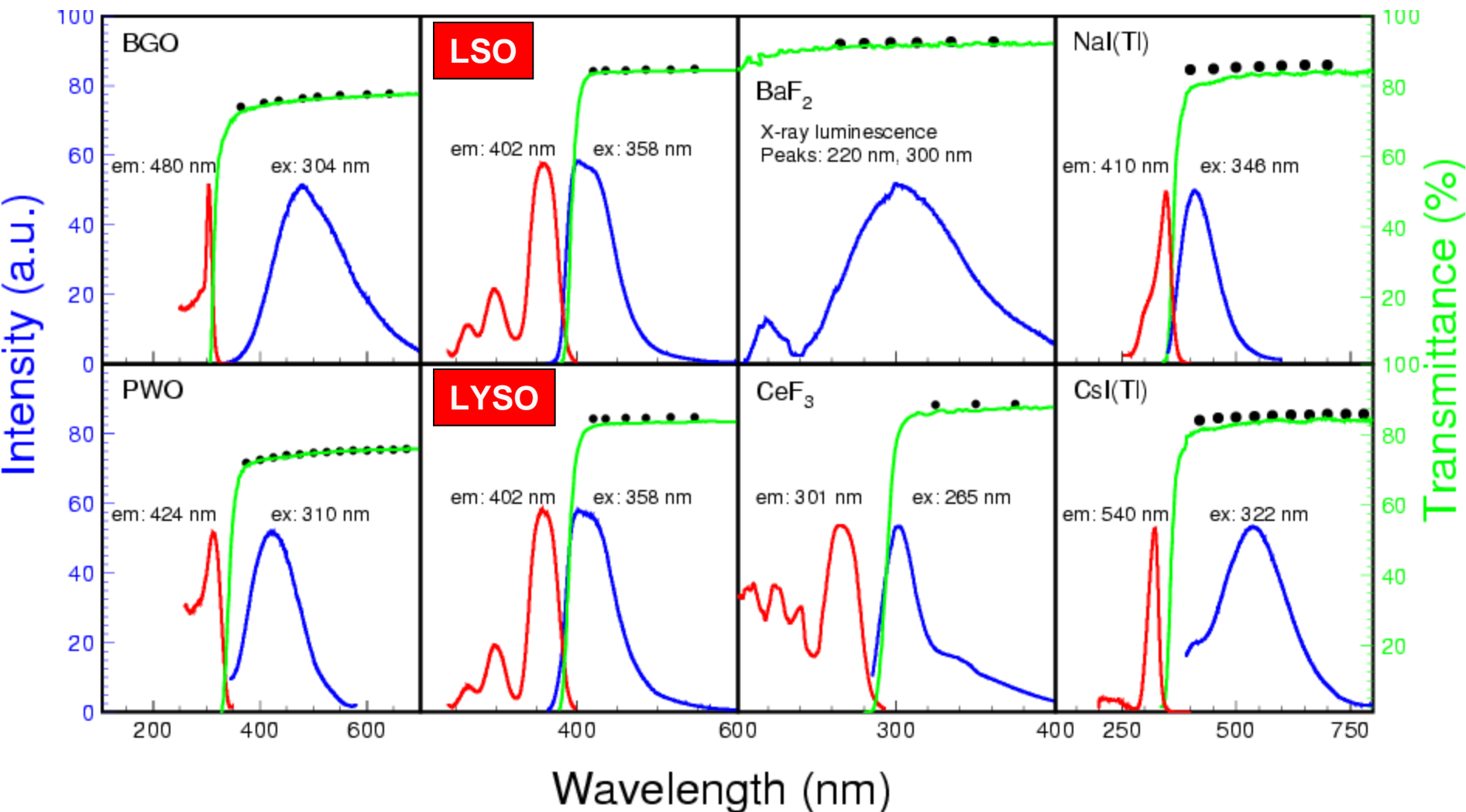
Excitation, Emission, Transmission



$$T_s = (1 - R)^2 + R^2(1 - R)^2 + \dots = (1 - R)/(1 + R), \text{ with}$$

$$R = \frac{(n_{\text{crystal}} - n_{\text{air}})^2}{(n_{\text{crystal}} + n_{\text{air}})^2}.$$

Black Dots: Theoretical limit of transmittance: NIM A333 (1993) 422





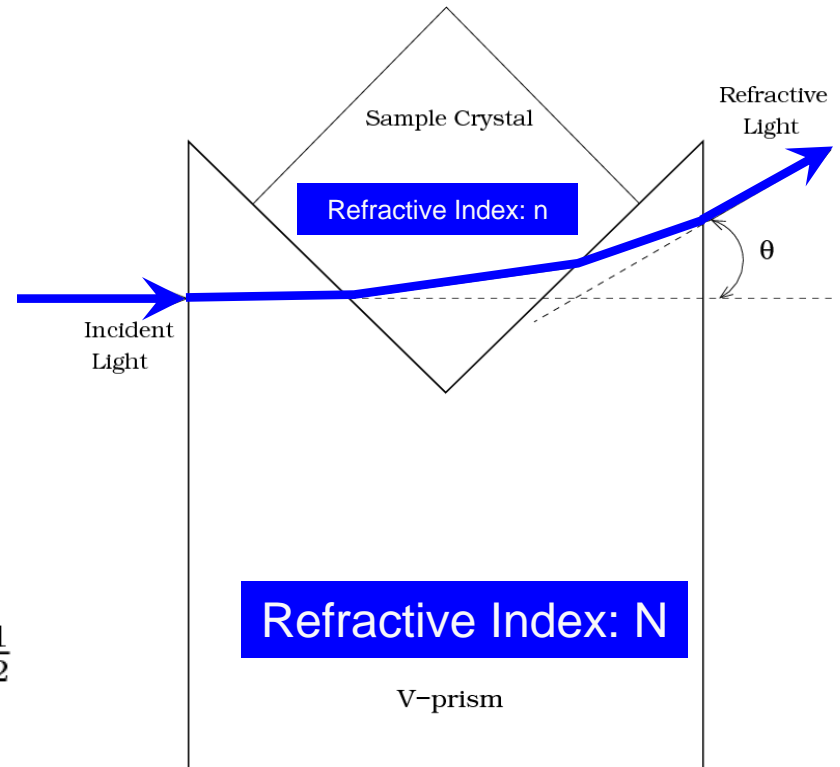
LSO/LYSO Refractive Index



Wavelength dependent measurement by a V-prism

- ◆ Cubic sample placed inside a V-prism
- ◆ Incident light shooting perpendicularly to one side of the prism
- ◆ The refractive index is calculated according to the following the equation:

$$n = (N^2 + \sin \theta \sqrt{N^2 - \sin^2 \theta})^{\frac{1}{2}}$$



λ (nm)	405	420	436	461	486	516	546
R. I.	1.833	1.827	1.822	1.818	1.813	1.810	1.806

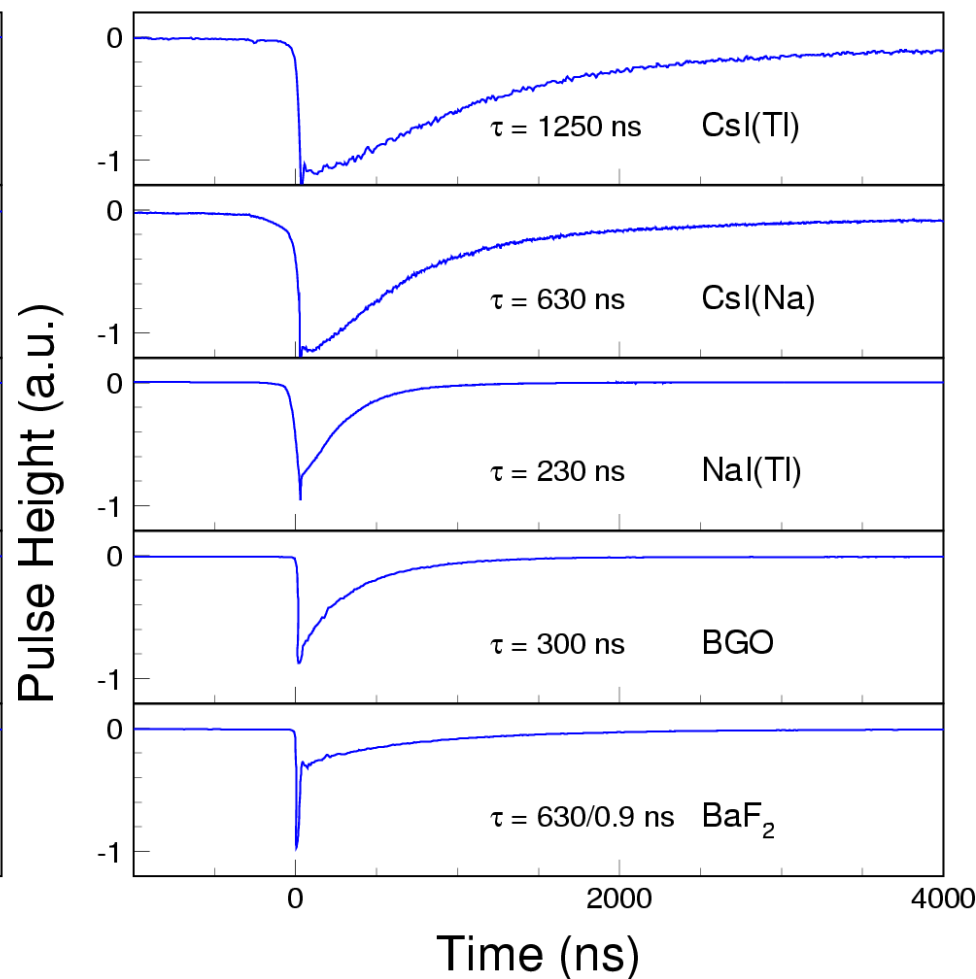
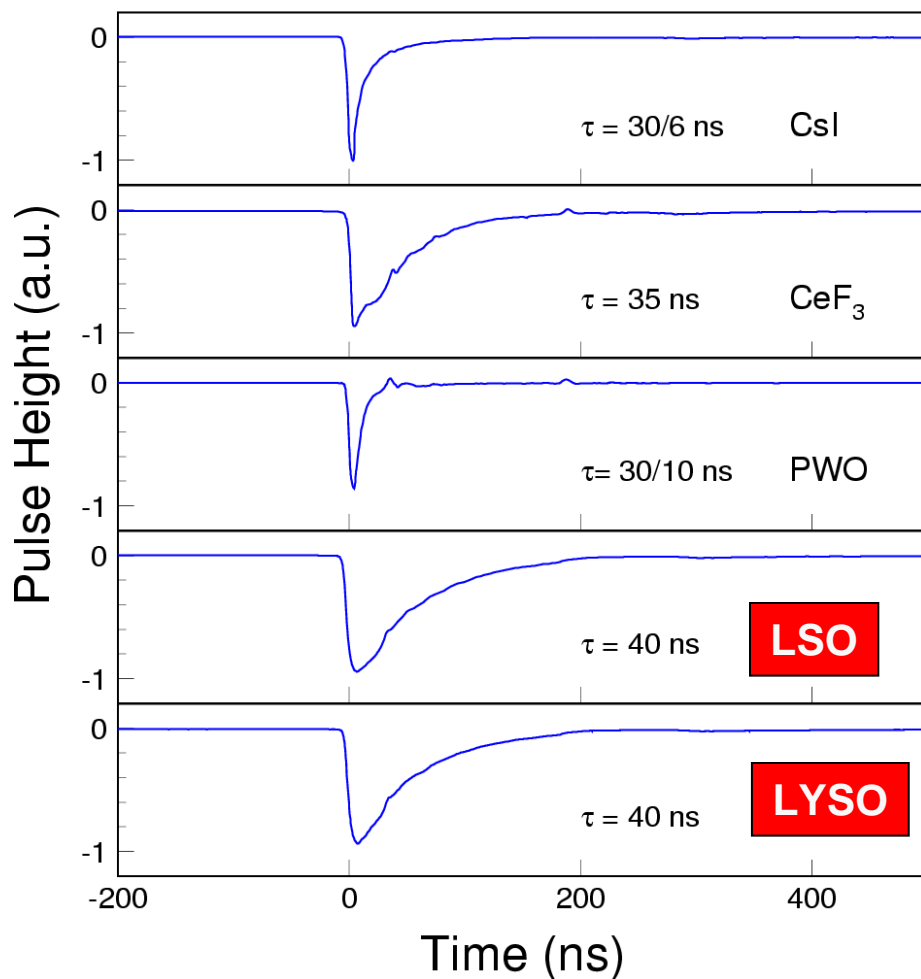
Scintillation Light Decay Time



Recorded with an Agilent 6052A digital scope

Fast Scintillators

Slow Scintillators

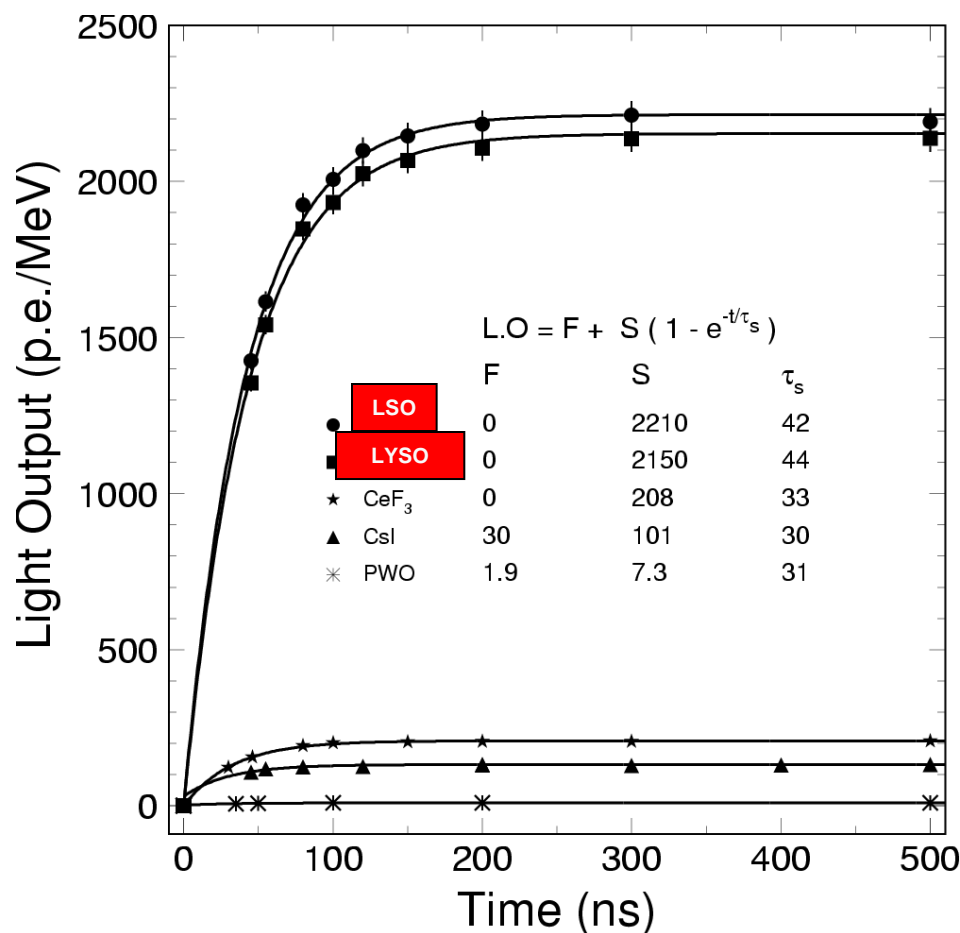


Light Output & Decay Kinetics

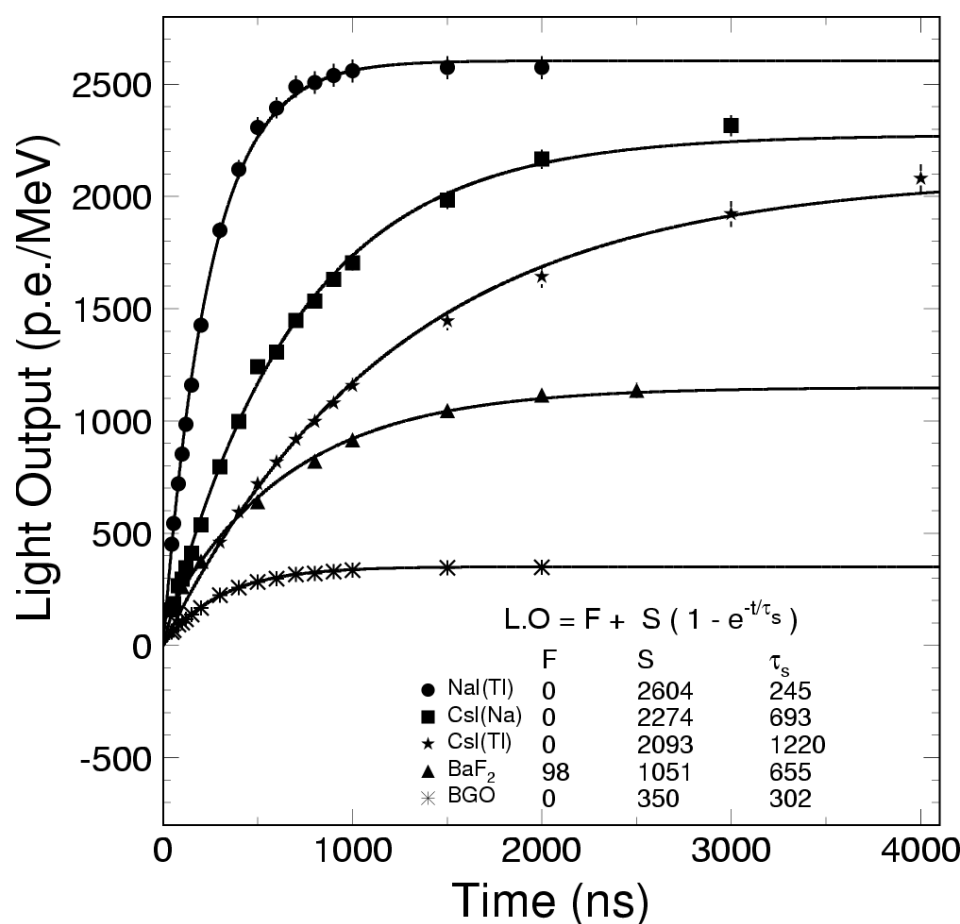


Measured with Philips XP2254B PMT (multi-alkali cathode)
p.e./MeV: LSO/LYSO is 6 & 230 times of BGO & PWO respectively

Fast Scintillators



Slow Scintillators

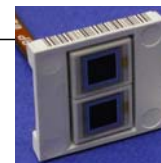
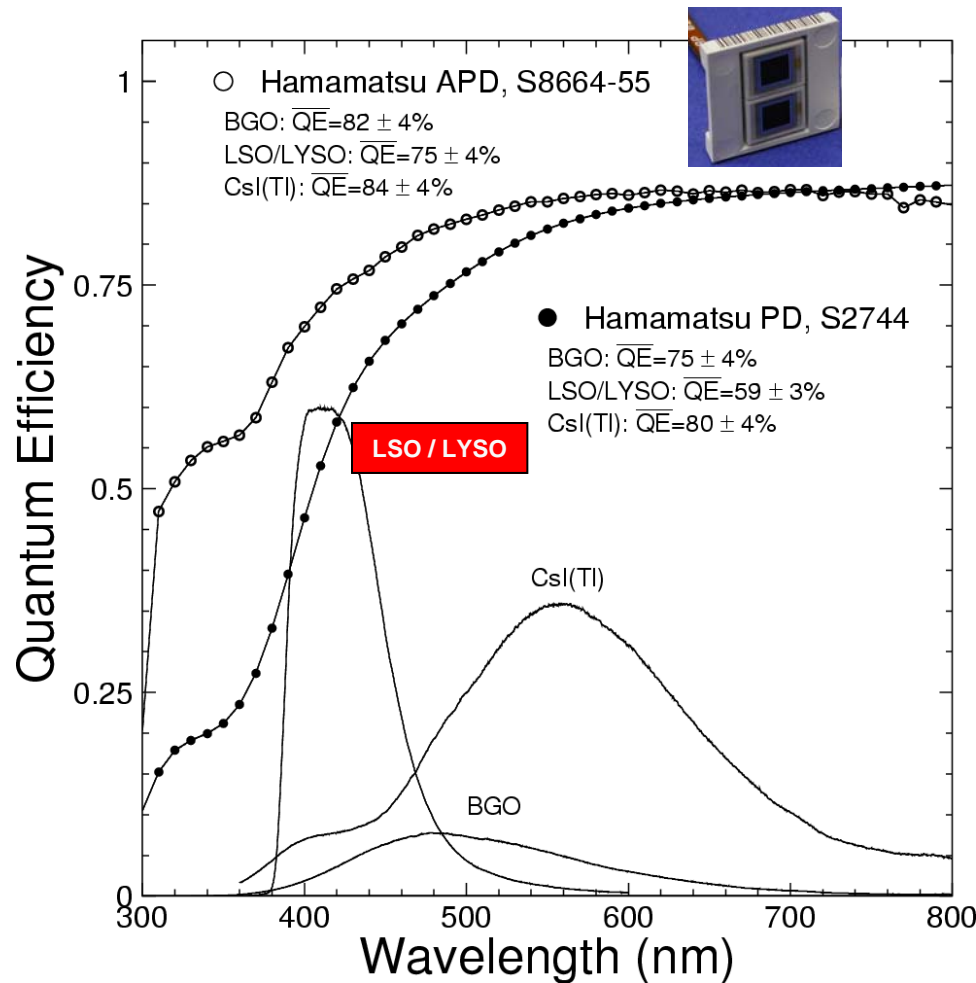
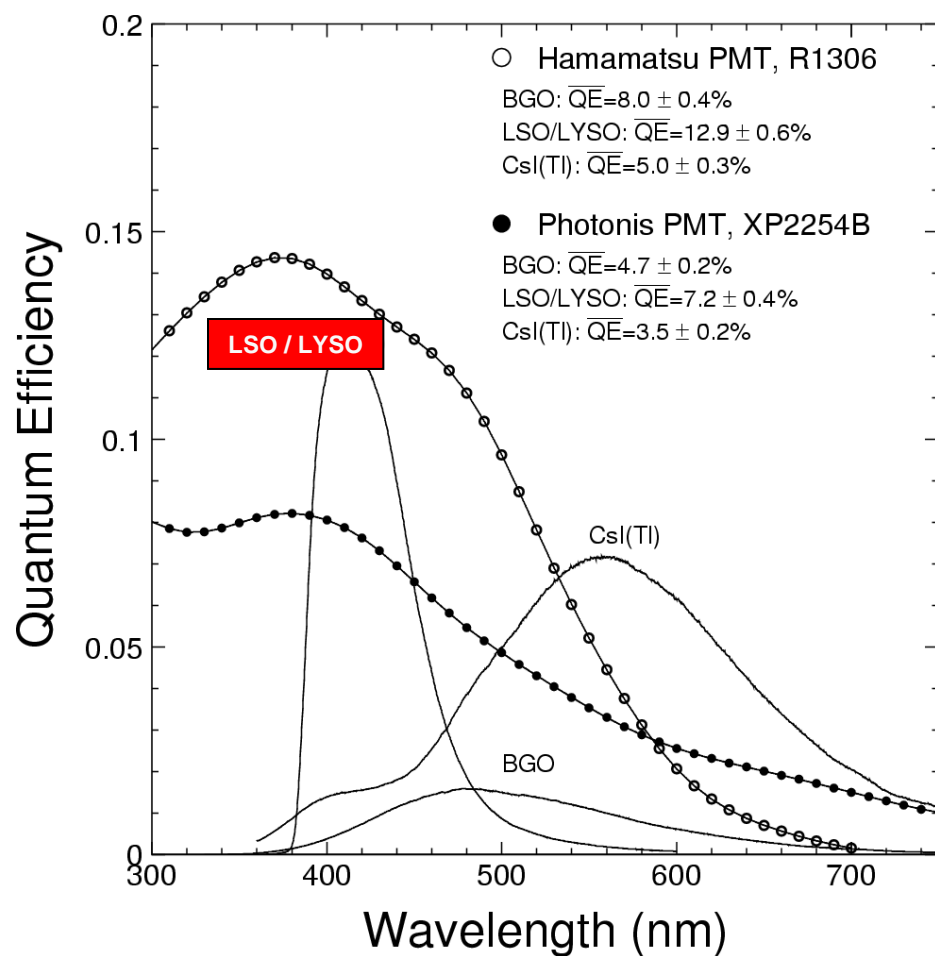




Emission Weighted Quantum Efficiency



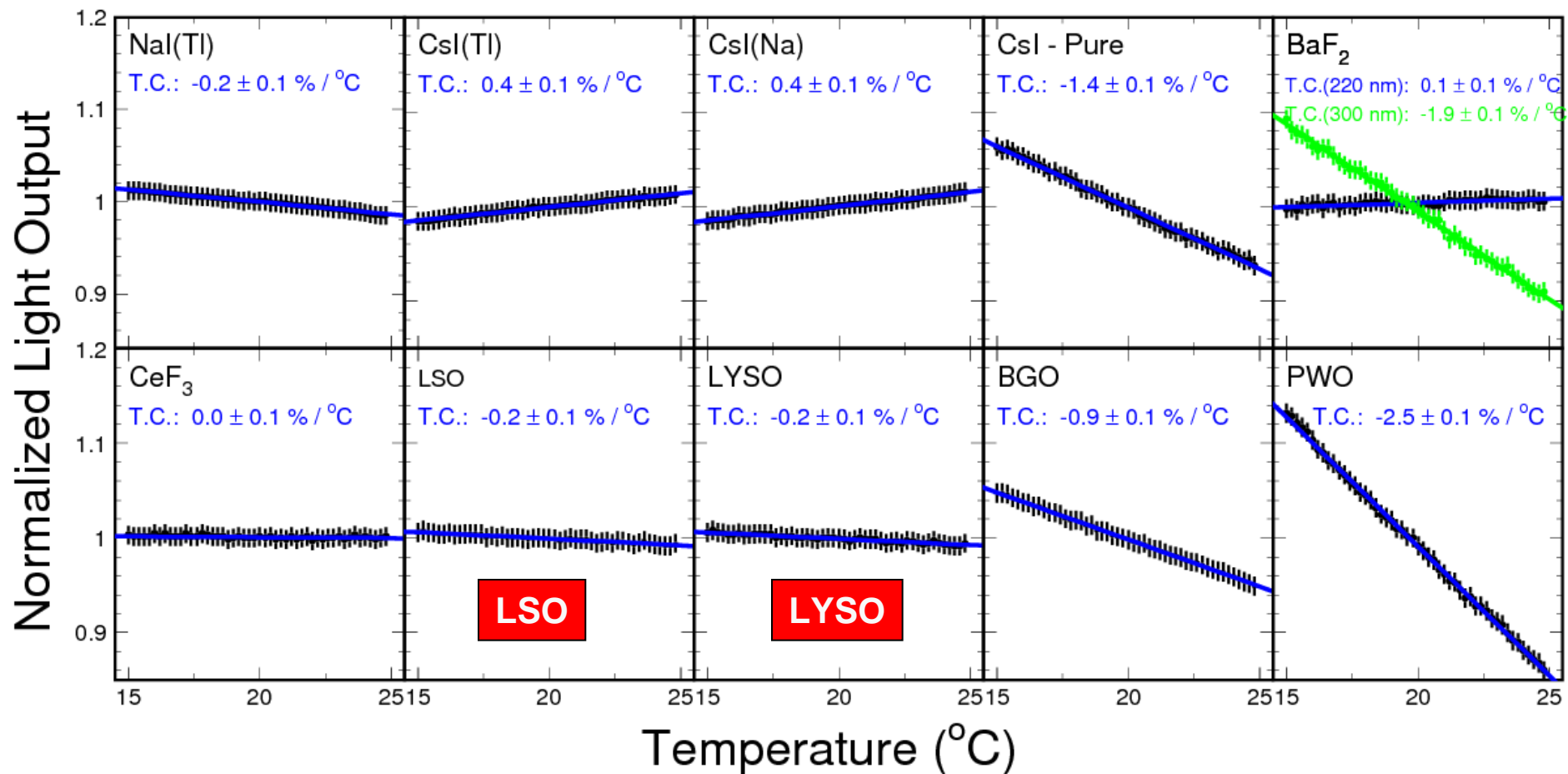
Taking out QE, L.O. of LSO/LYSO is 4/200 times BGO/PWO
Hamamatsu S8664-55 APD has QE 75% for LSO/LYSO



Light Output Temperature Coefficient



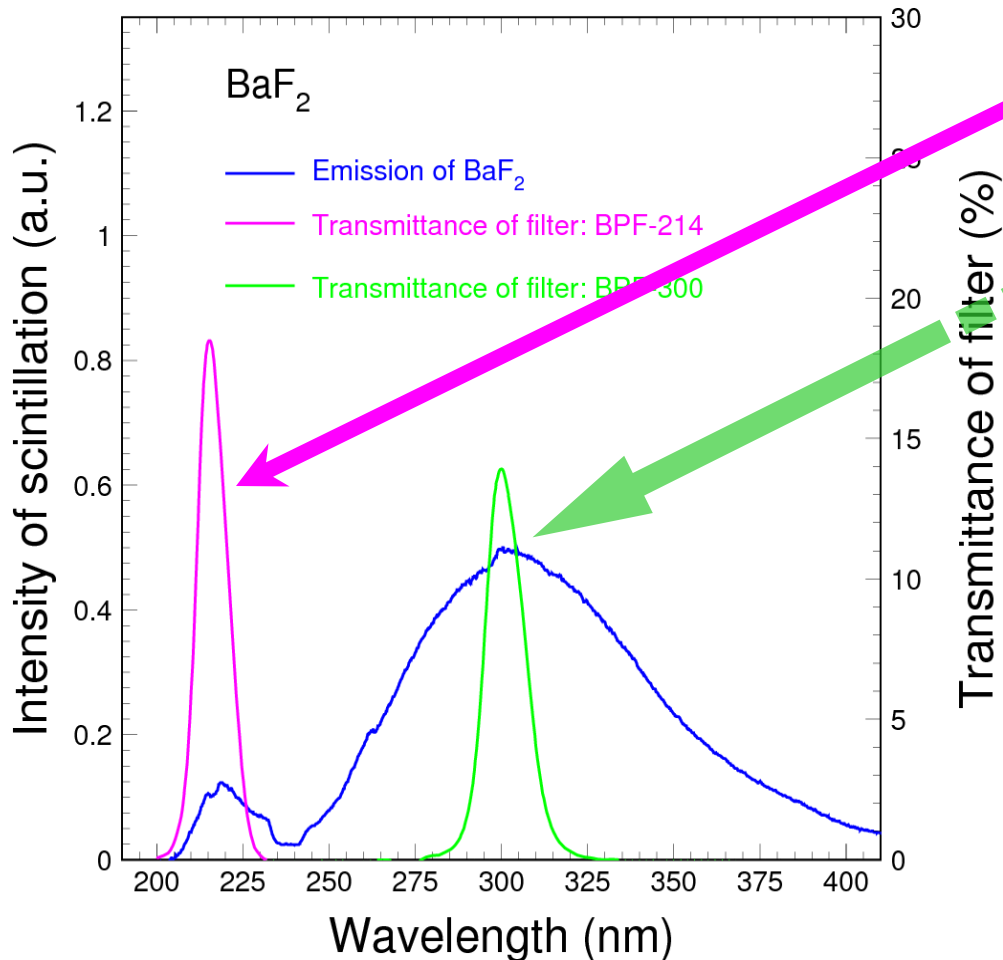
Temperature Range: 15°C ~ 25°C



BaF₂: Fast and Slow Components



Two filters used to select scintillation component



Transmittance for filter BPF-214
(fast component)

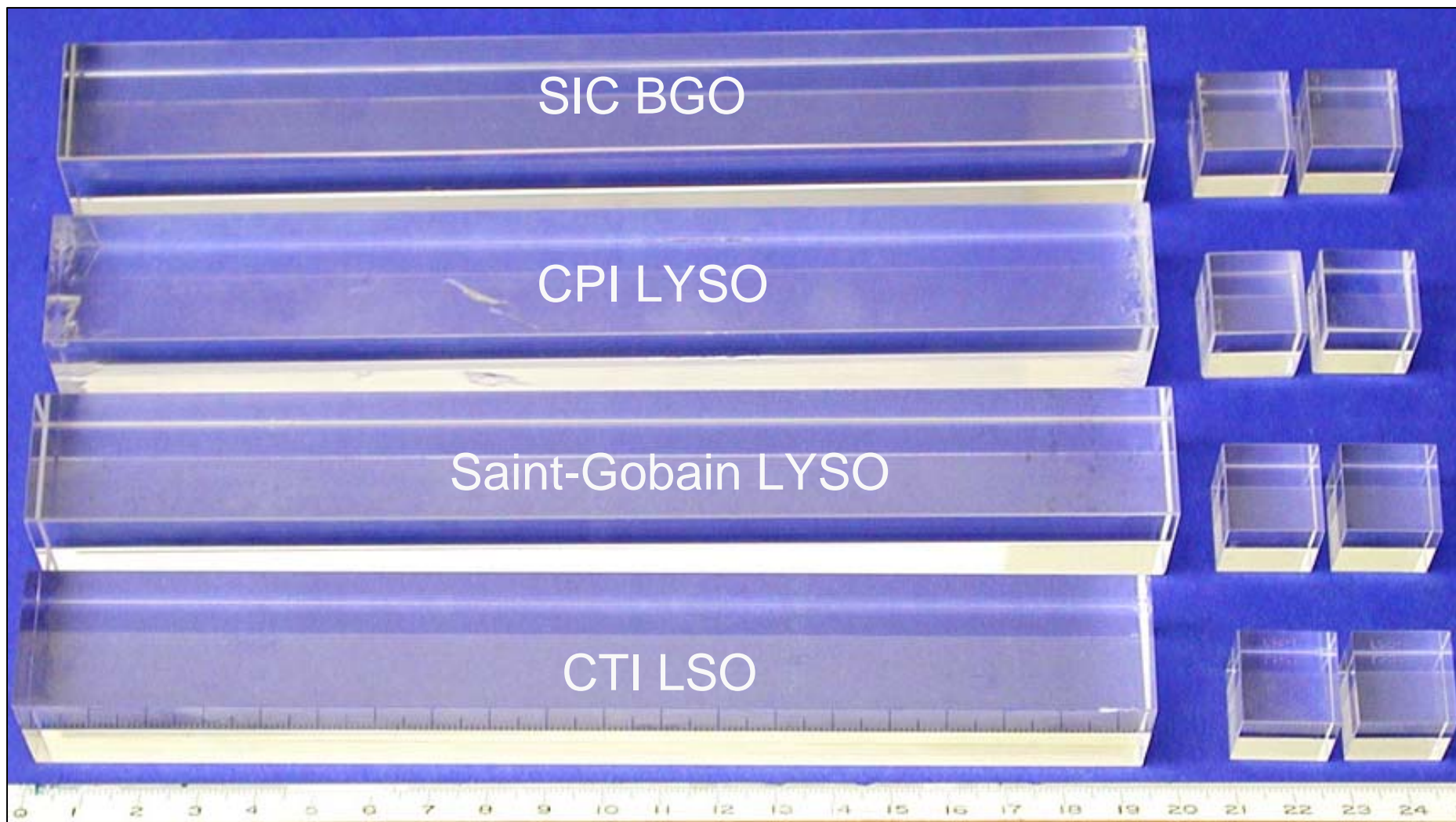
Transmittance for filter BPF-300
(slow component)

- Scintillation of BaF₂ has two components: the fast one peaked at 220 nm while the slow one peaked at 300 nm.
- Special band pass filters were used to measure the light output temperature coefficients for individual component.



Comparison of BGO, LSO & LYSO

2.5 x 2.5 x 20 cm (18 X_0) Bar



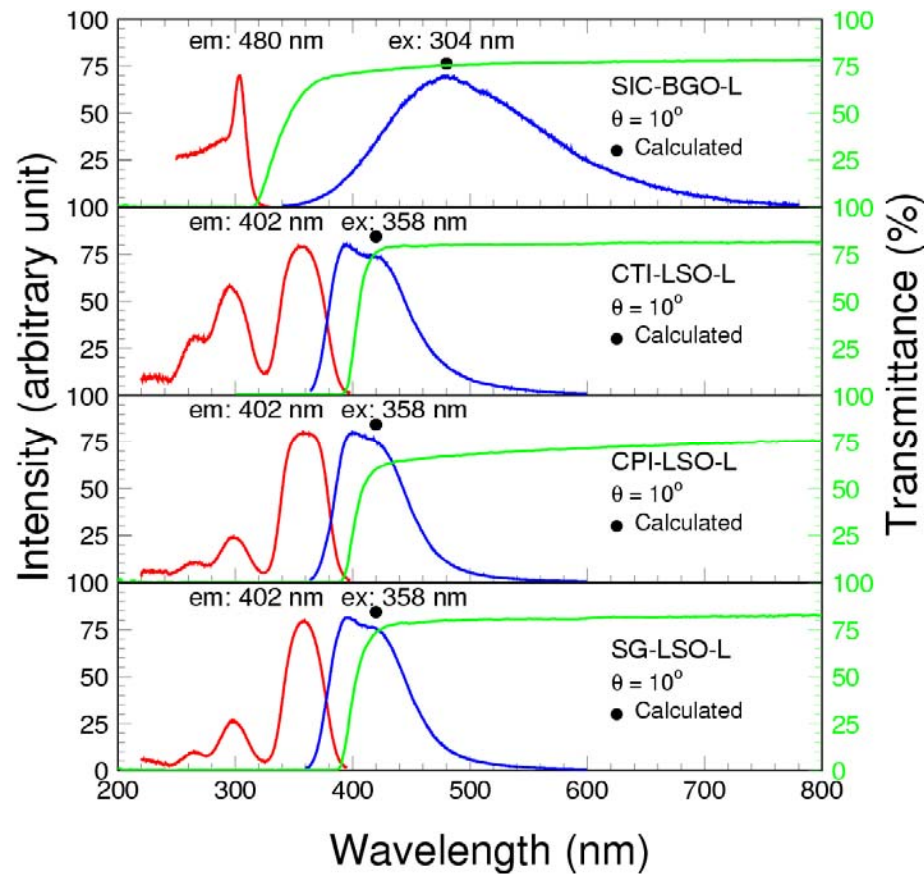
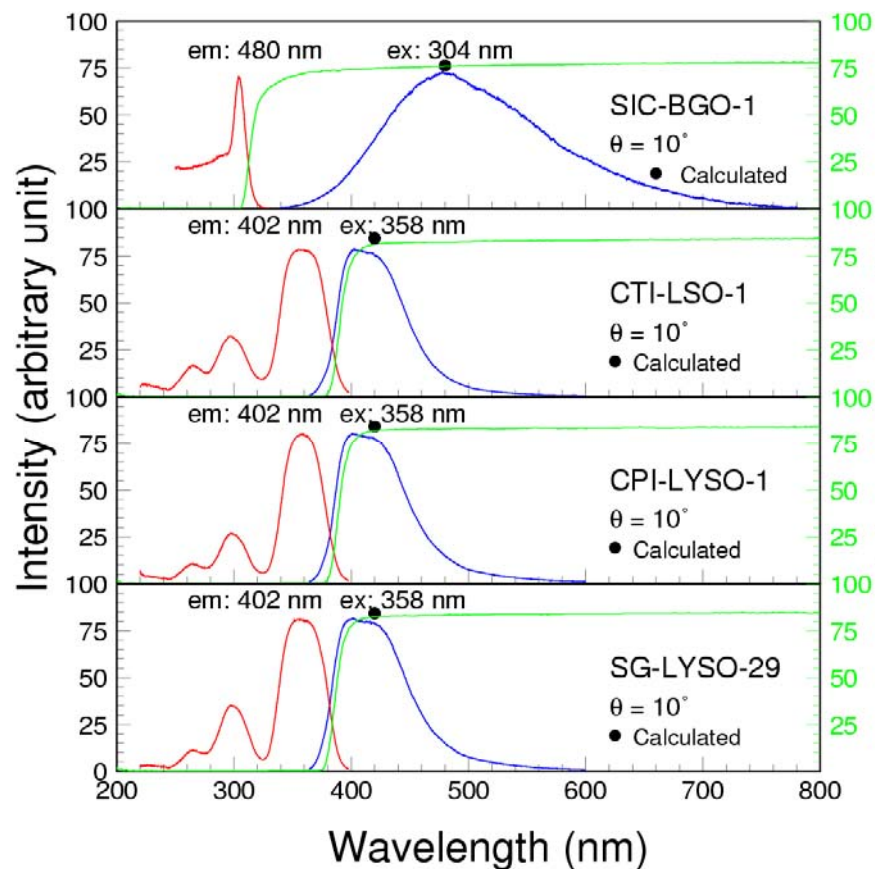
Excitation, Emission, Transmission



Identical transmittance, emission & excitation spectra
Part of emitted light may be self-absorbed in long samples

1.7 cm Cube

2.5 x 2.5 x 20 cm Bar

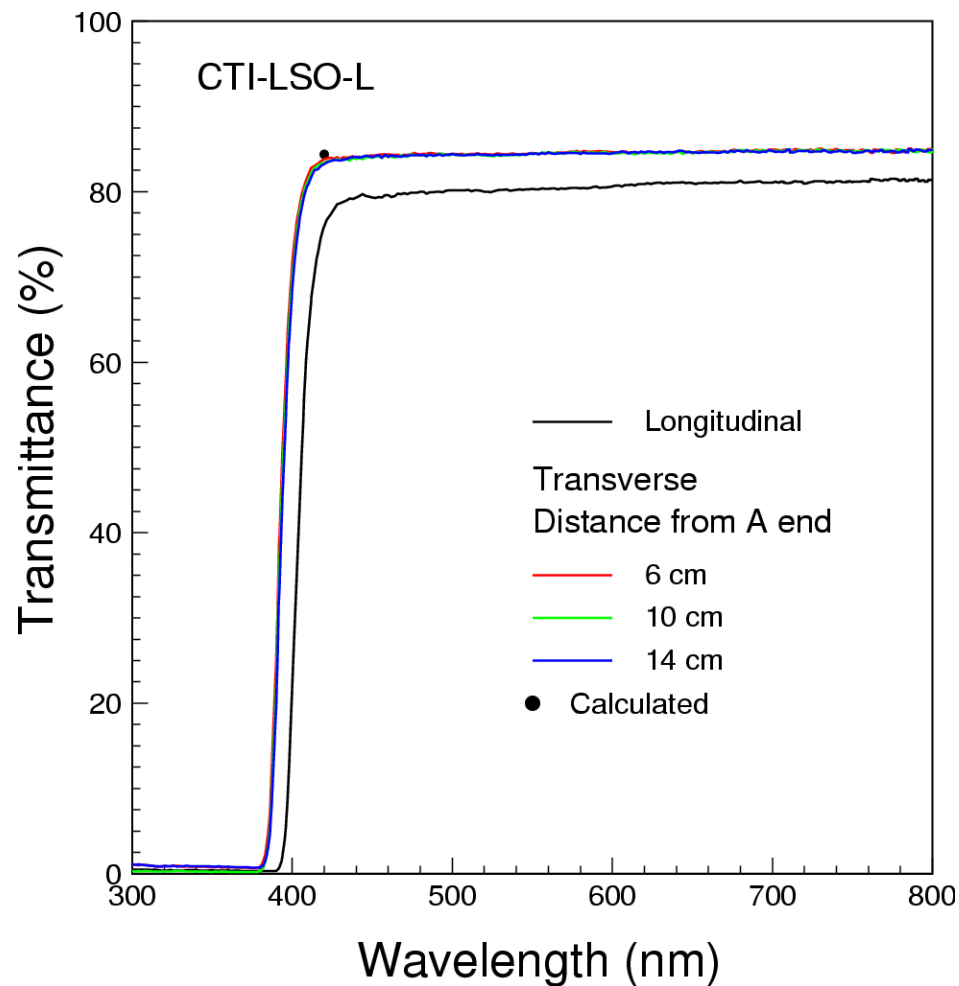
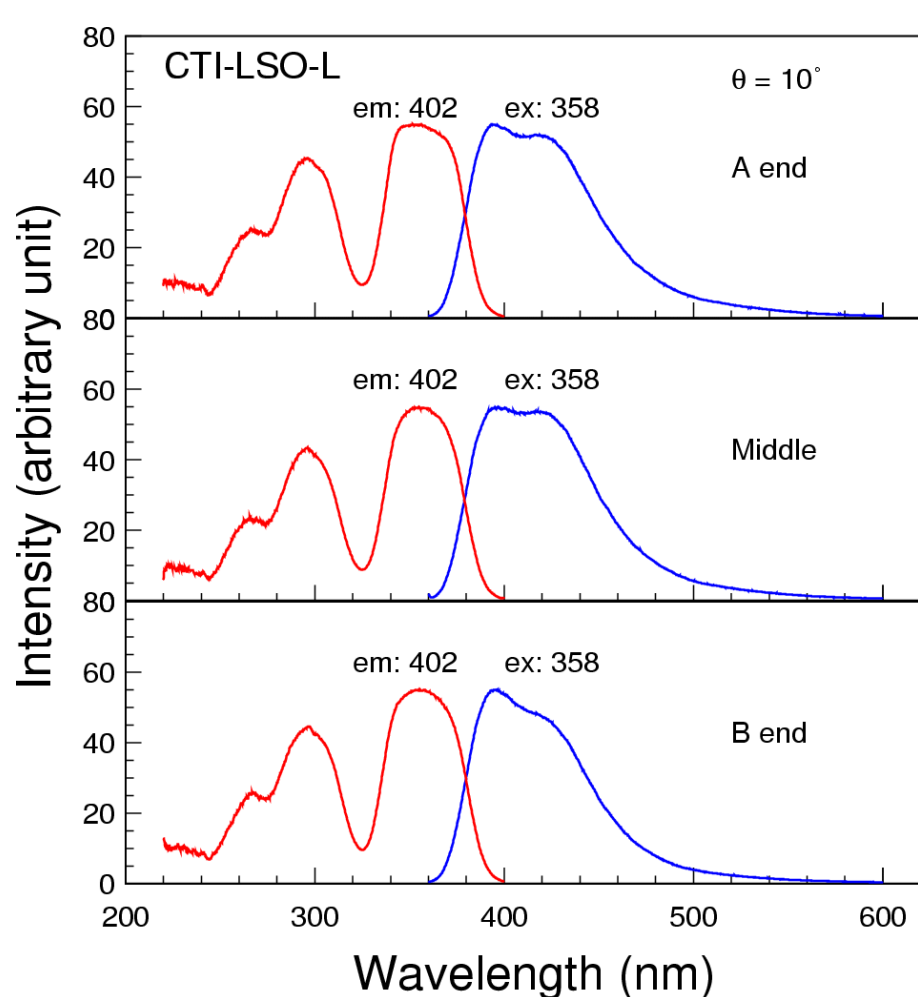




CTI LSO: Longitudinal Uniformity



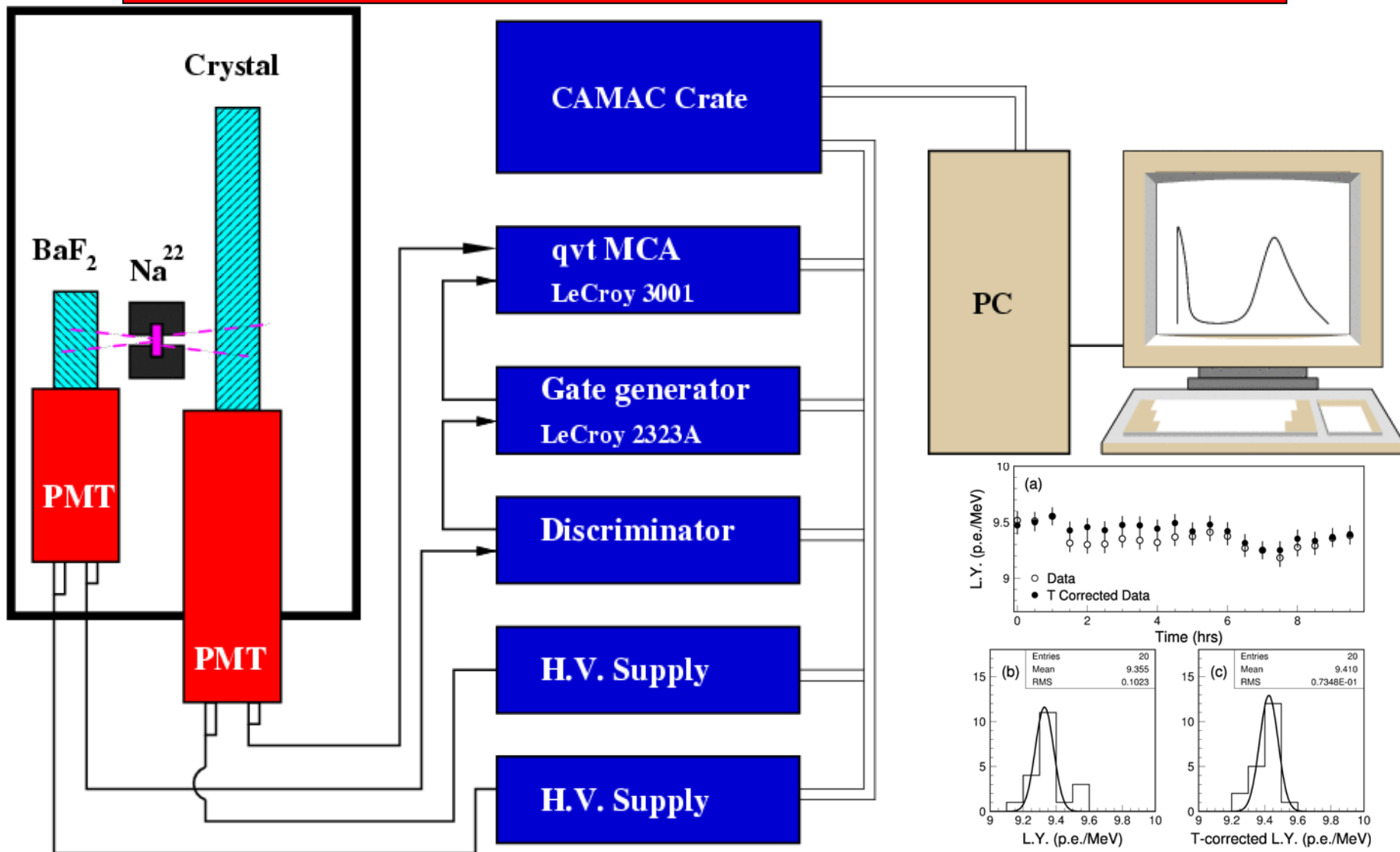
No longitudinal variation in optical properties
Transverse transmittance approaches theoretical limit



PMT Based Readout with Coincidence



Systematic error with repeated mounts & measurements: $< 1\%$





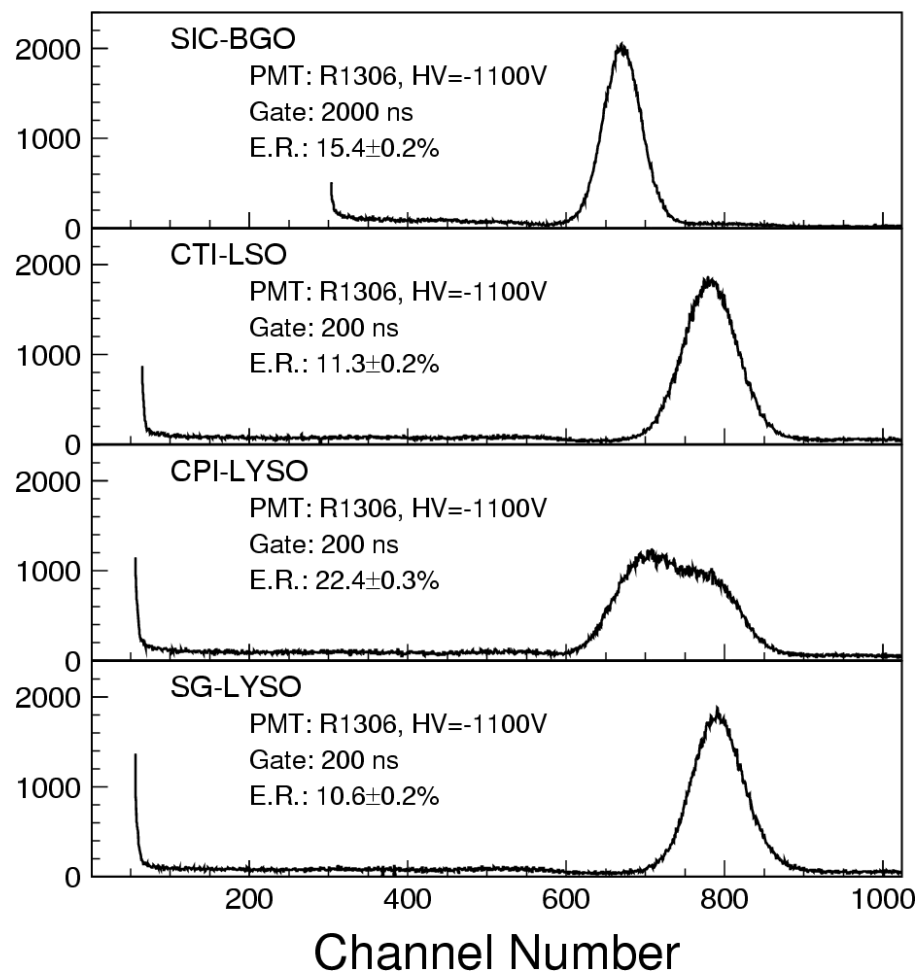
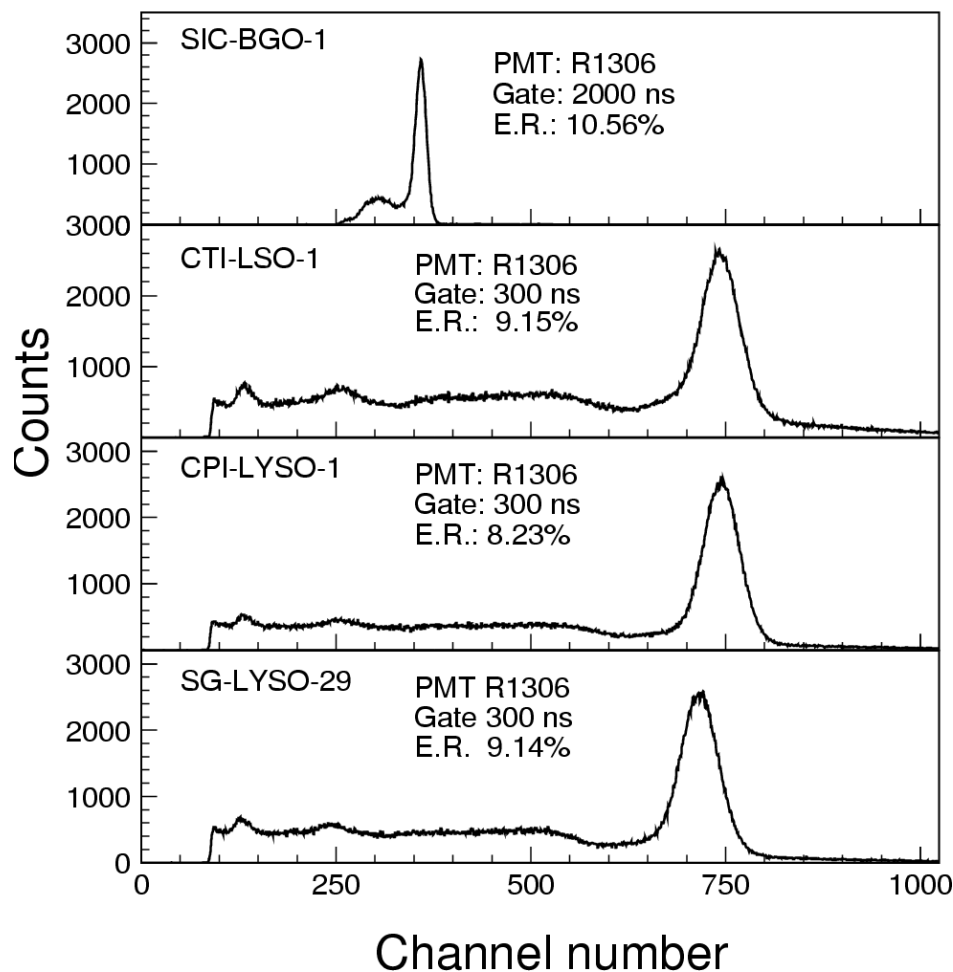
LSO/LYSO Resolution with PMT



~10% FWHM resolution for ^{22}Na source (0.51 MeV)

1.7 cm Cube

2.5 x 2.5 x 20 cm Bar



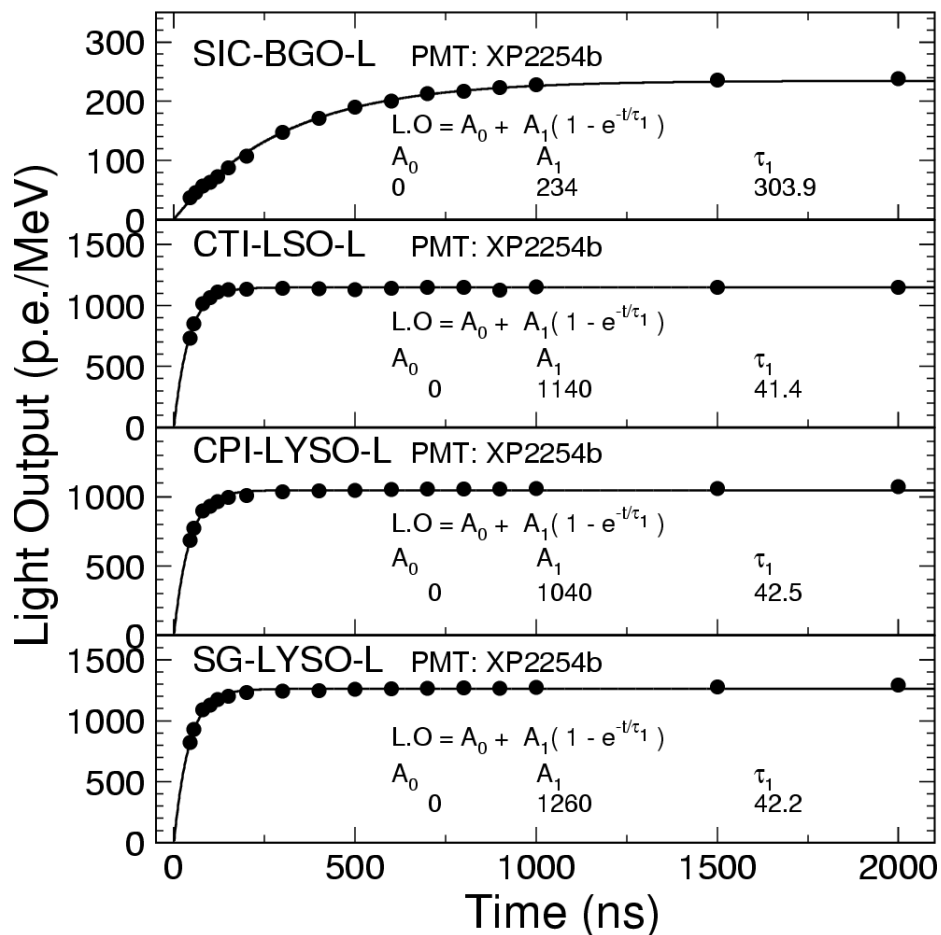
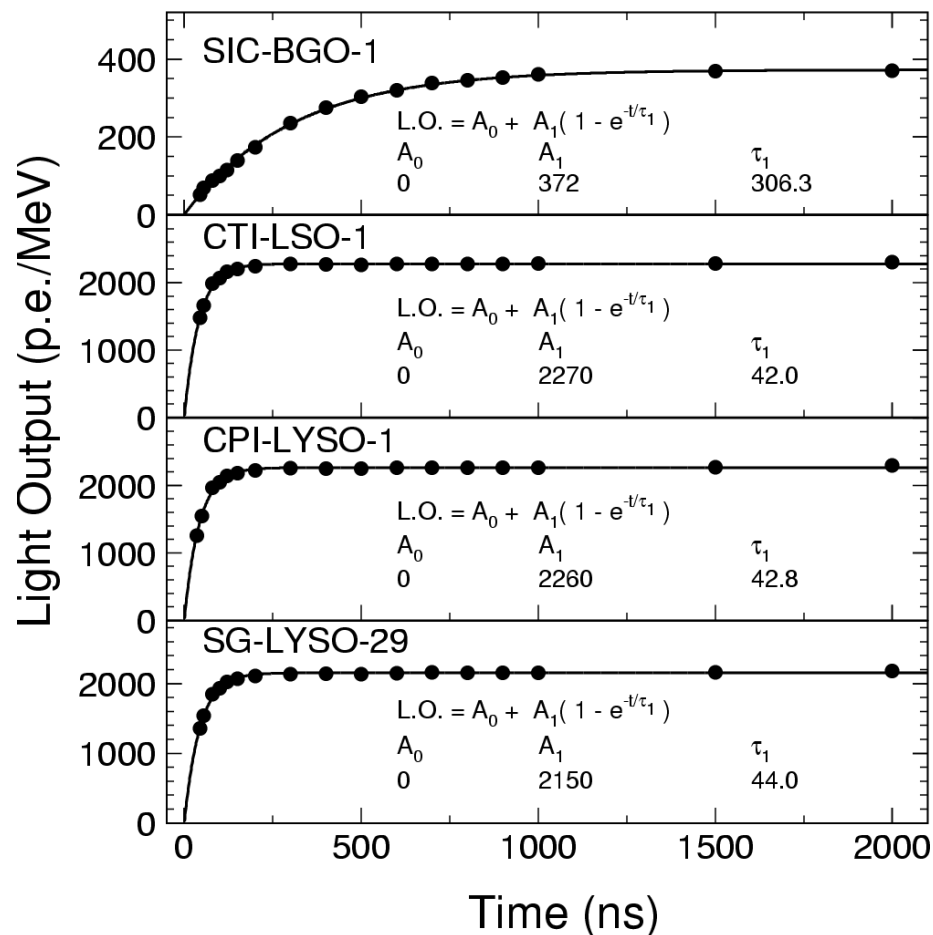
LSO/LYSO Light Output with PMT



1,200 p.e./MeV, 5/230 times of BGO/PWO

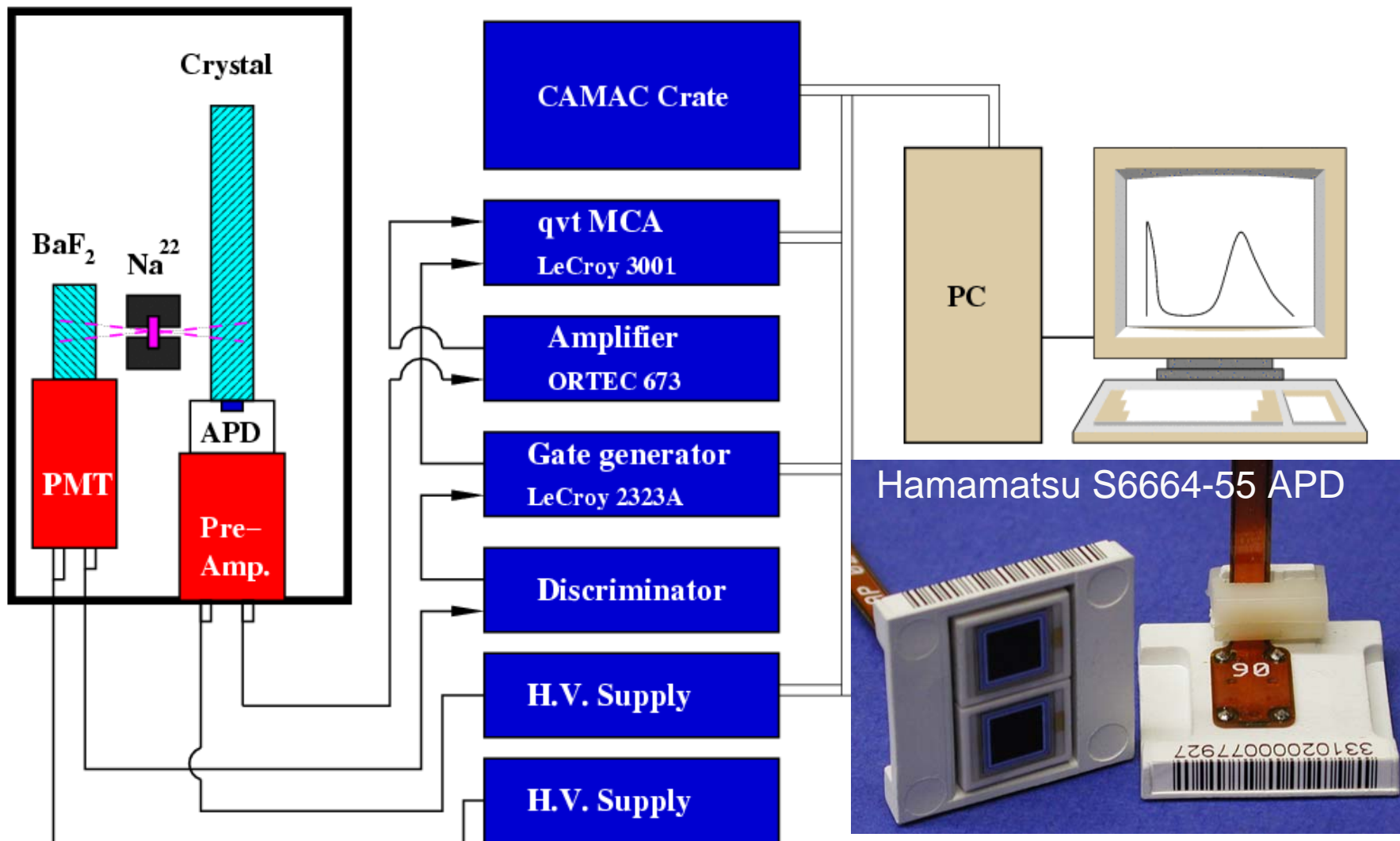
1.7 cm Cube

2.5 x 2.5 x 20 cm Bar



APD Based Readout with Coincidence

Two Hamamatsu S6664-55 APD, Canberra 2003 BT preamplifier and ORTEC 673 shaping amplifier with shaping time 250 ns

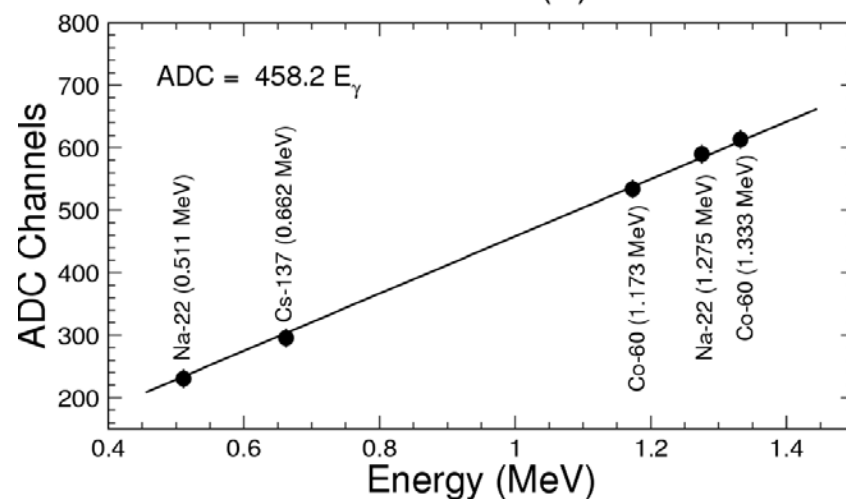
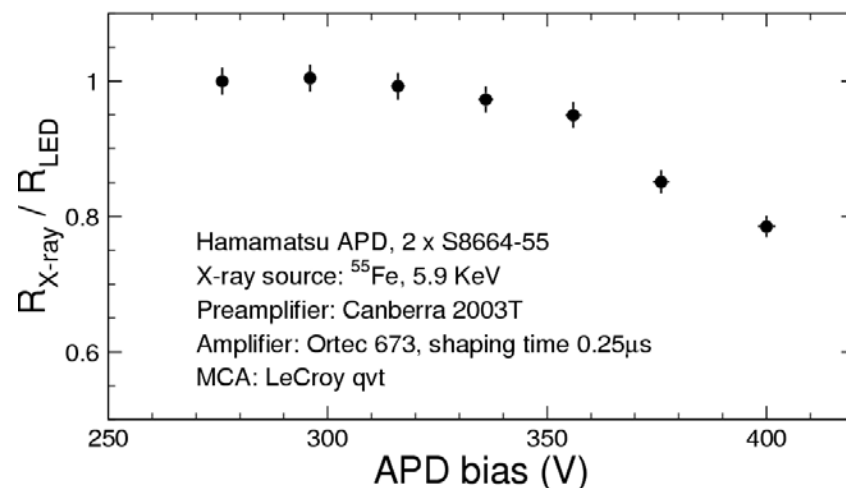
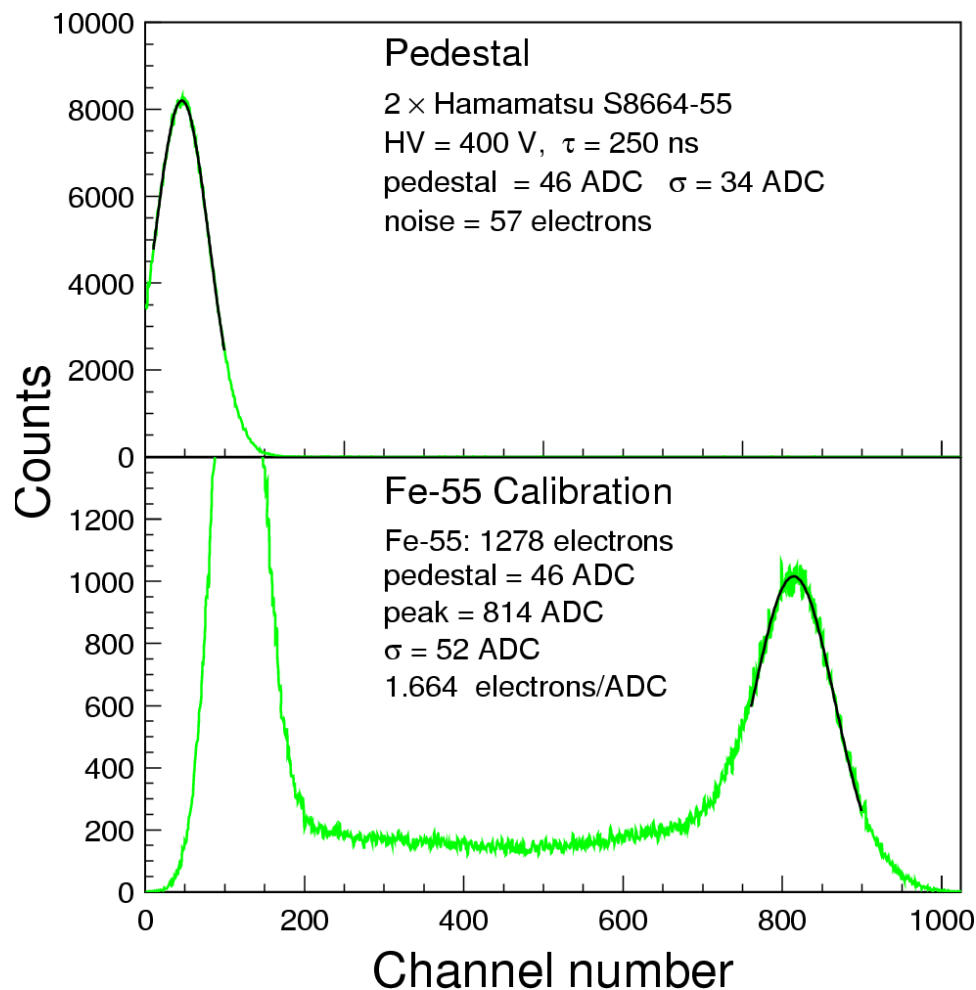




Calibration of the APD Readout



Pedestal: 34 ADC, corresponding to 57 electrons
Corrections for 5.9 keV X-ray: 78%; Good linearity



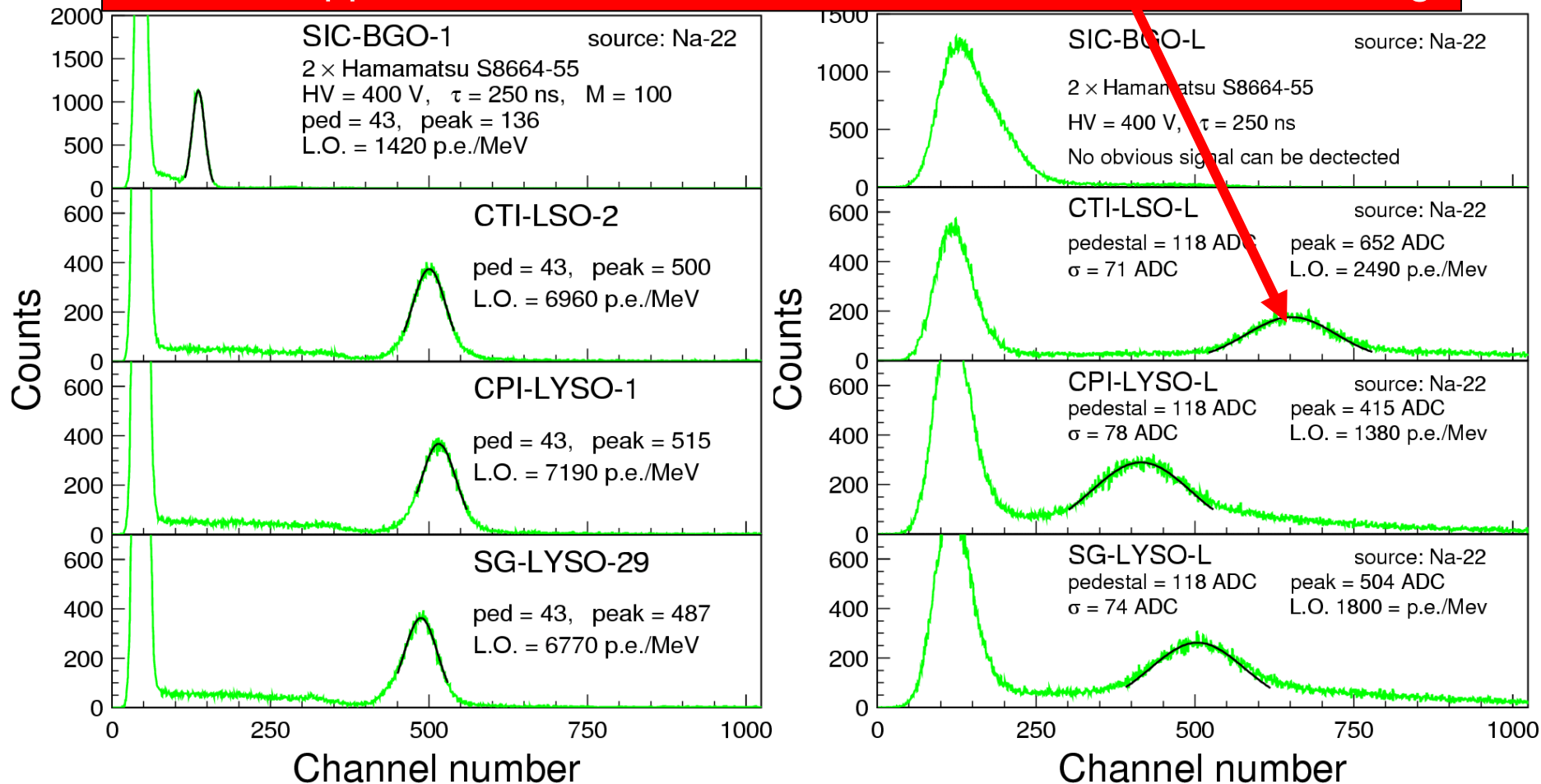


LSO/LYSO Light Output with APD



1,500 p.e./MeV, 4/200 times of BGO/PWO, Noise < 40 keV

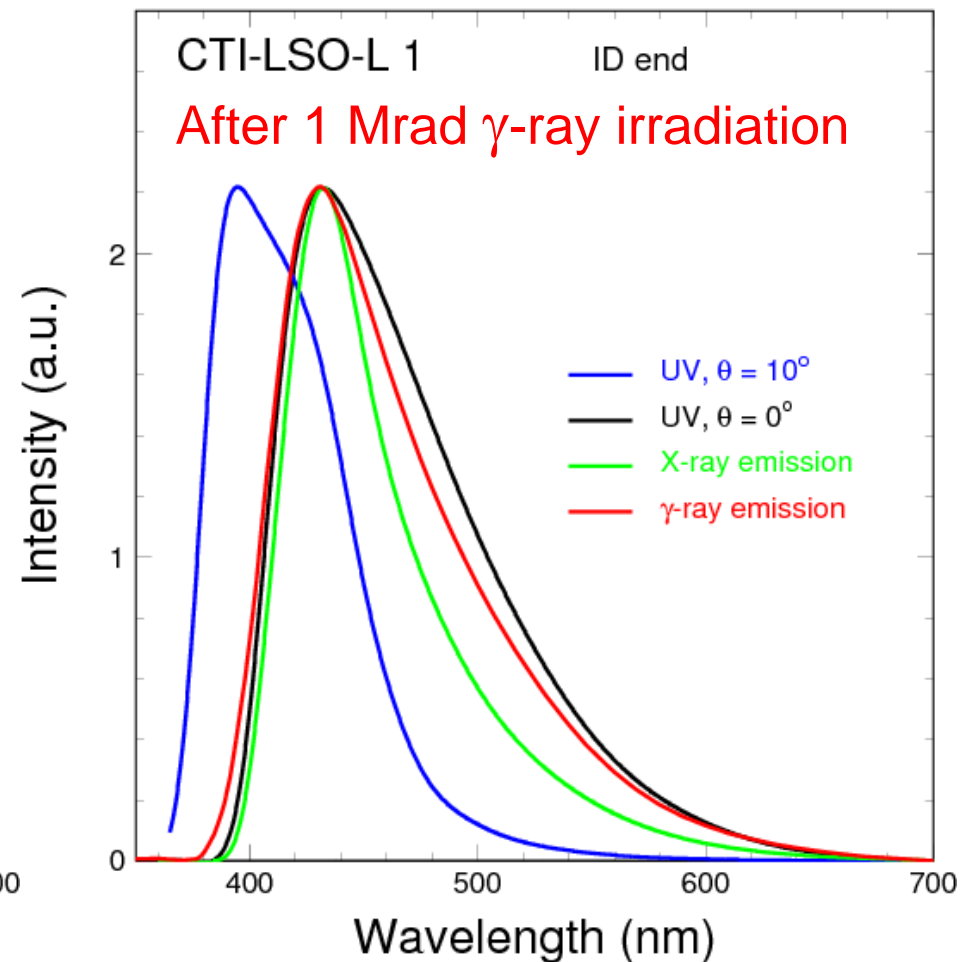
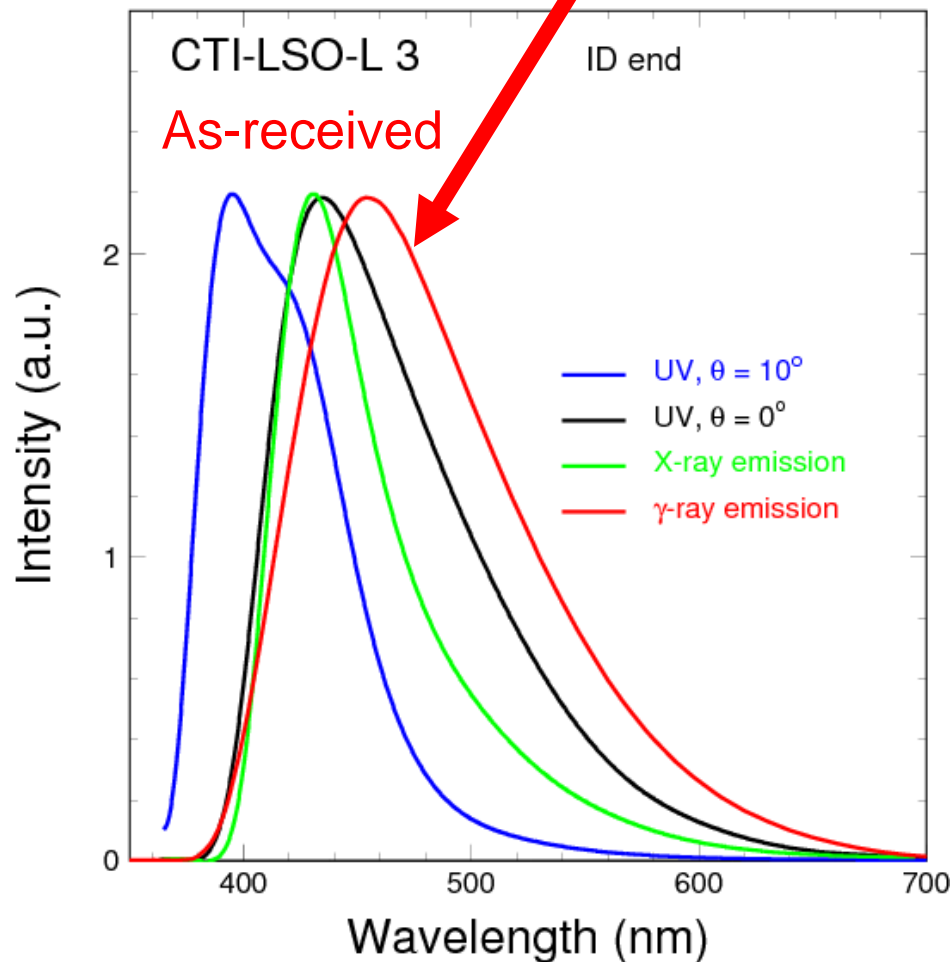
Discrepancy reported in NSS05: LSO has more light output, which disappeared after irradiation to 1 Mrad and thermal annealing.



LSO Emission Spectra

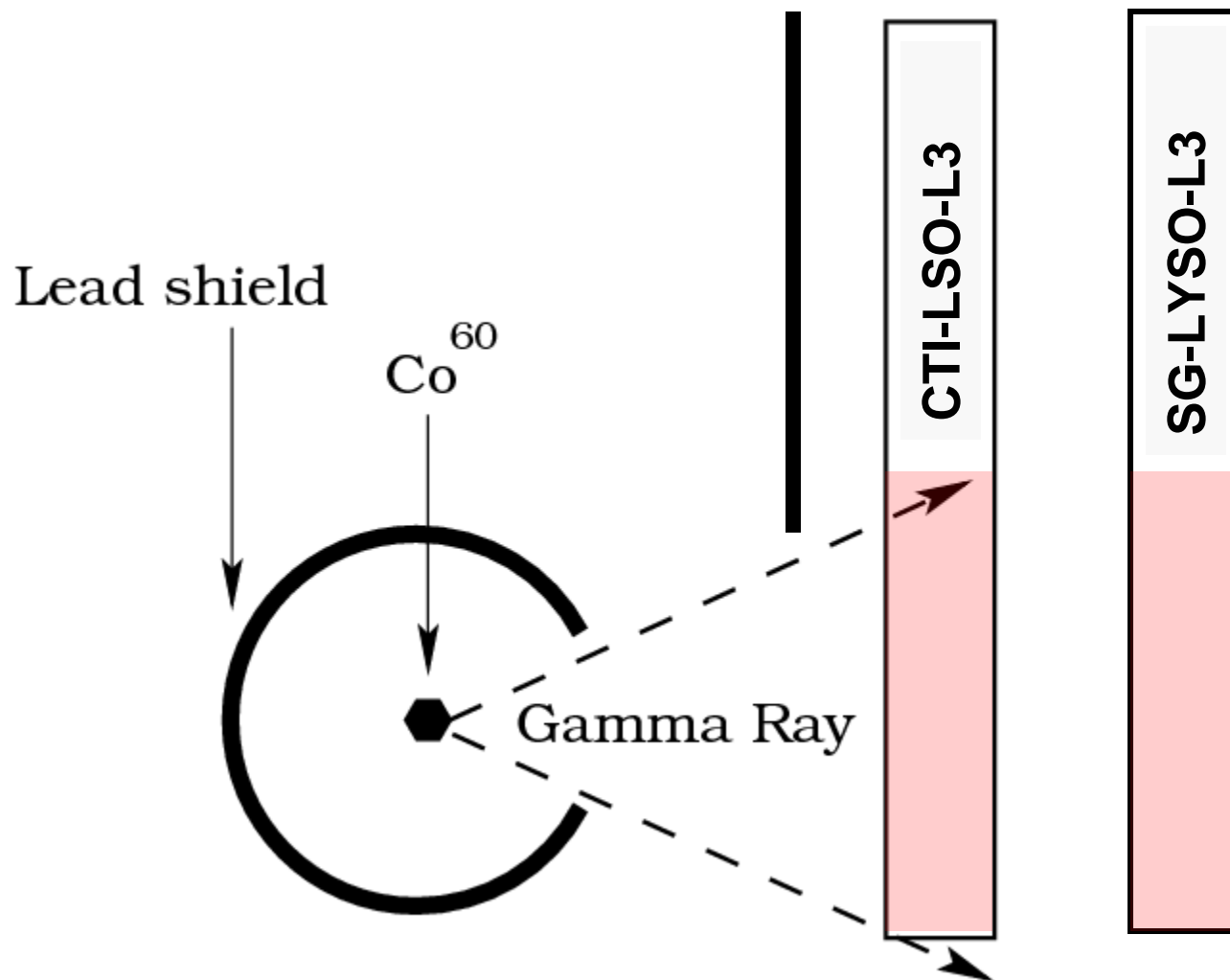


All emission spectra are similar to that of LYSO, except that γ -ray excited emission has a “red shift”, which disappeared after irradiations with γ -ray.





γ -Ray Irradiation on Sample's ID End

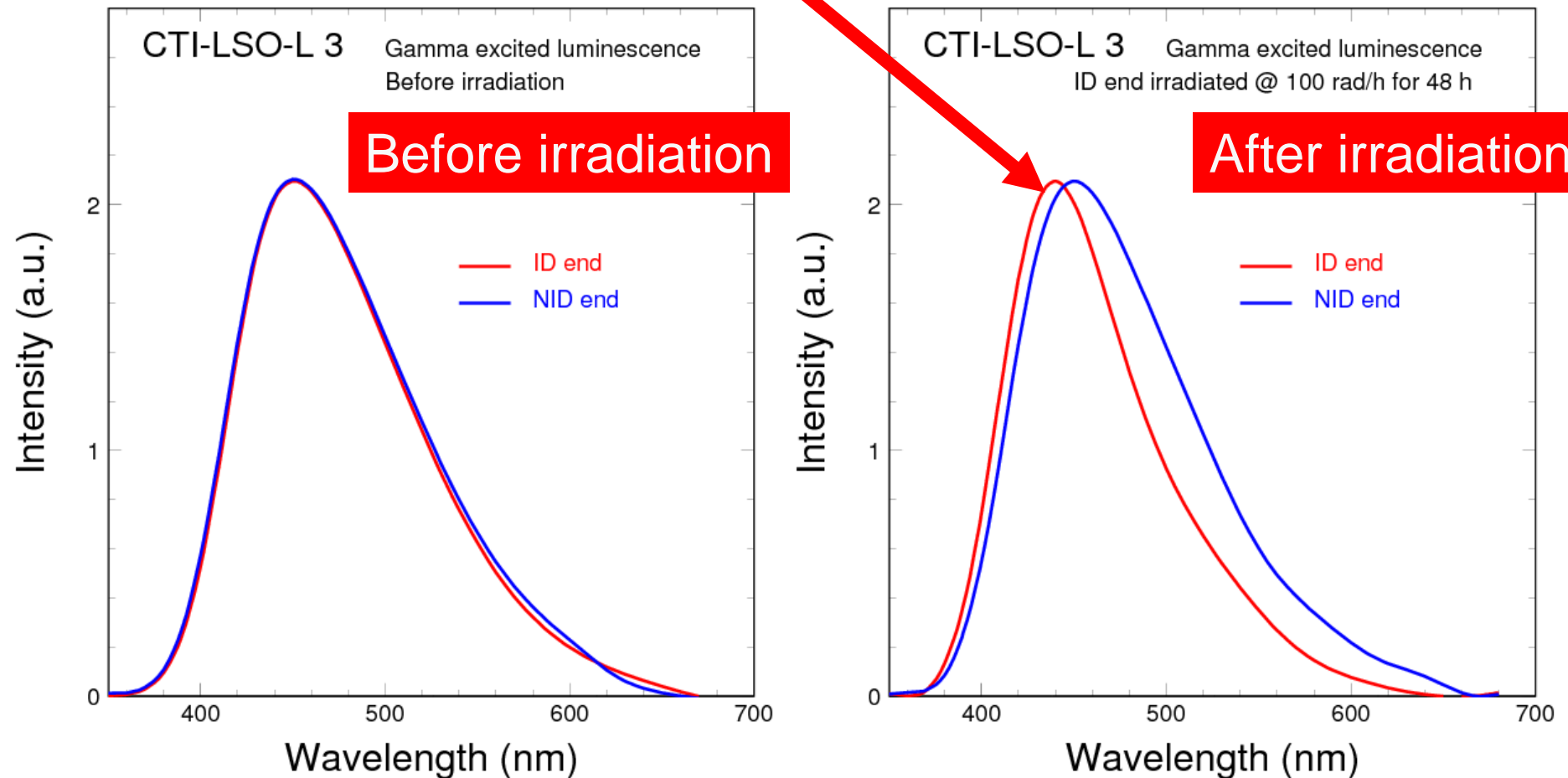


ID End received
~5,000 rad



LSO: γ -Ray Excited Emission Spectra

The emission peak of sample's irradiated ID end has a ~ 15 nm “blue” shift

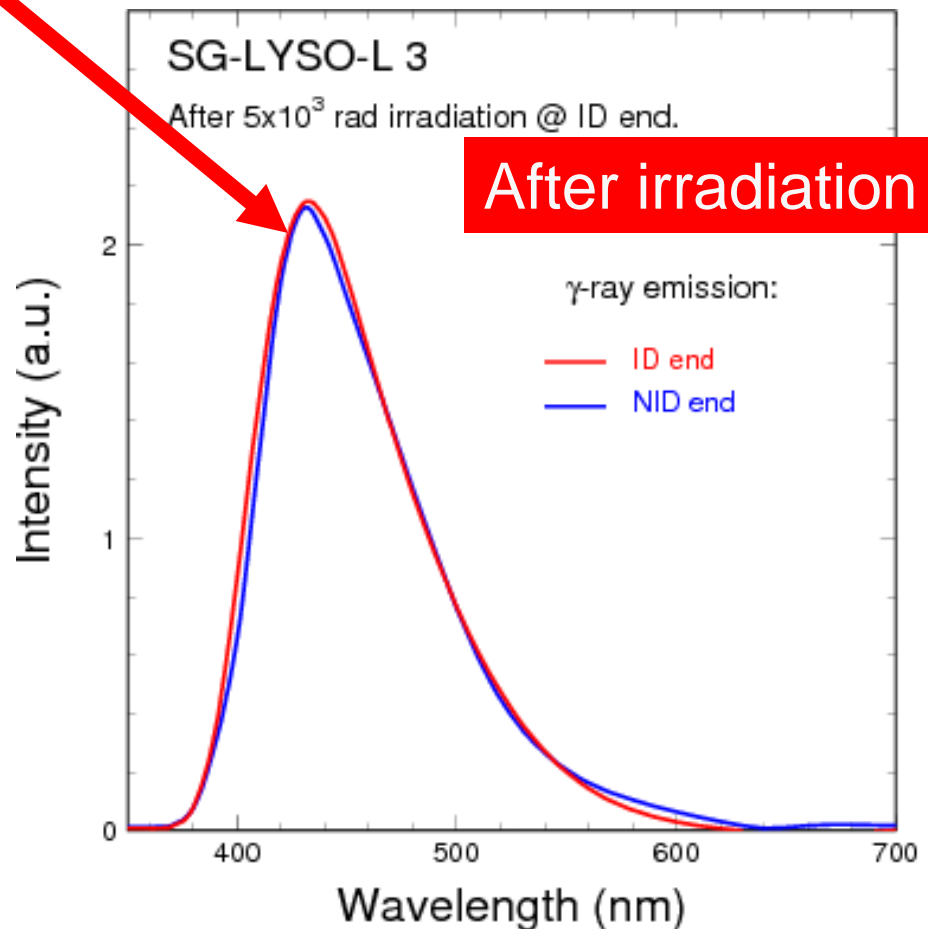
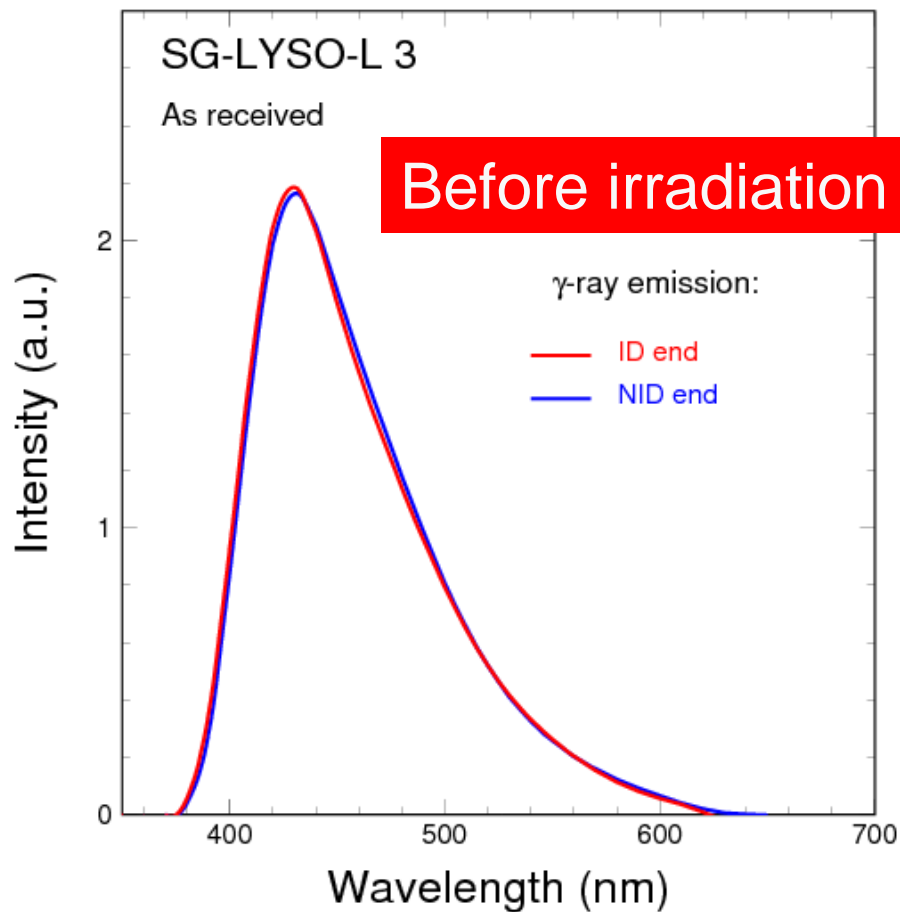




LYSO: γ -Ray Excited Emission Spectra



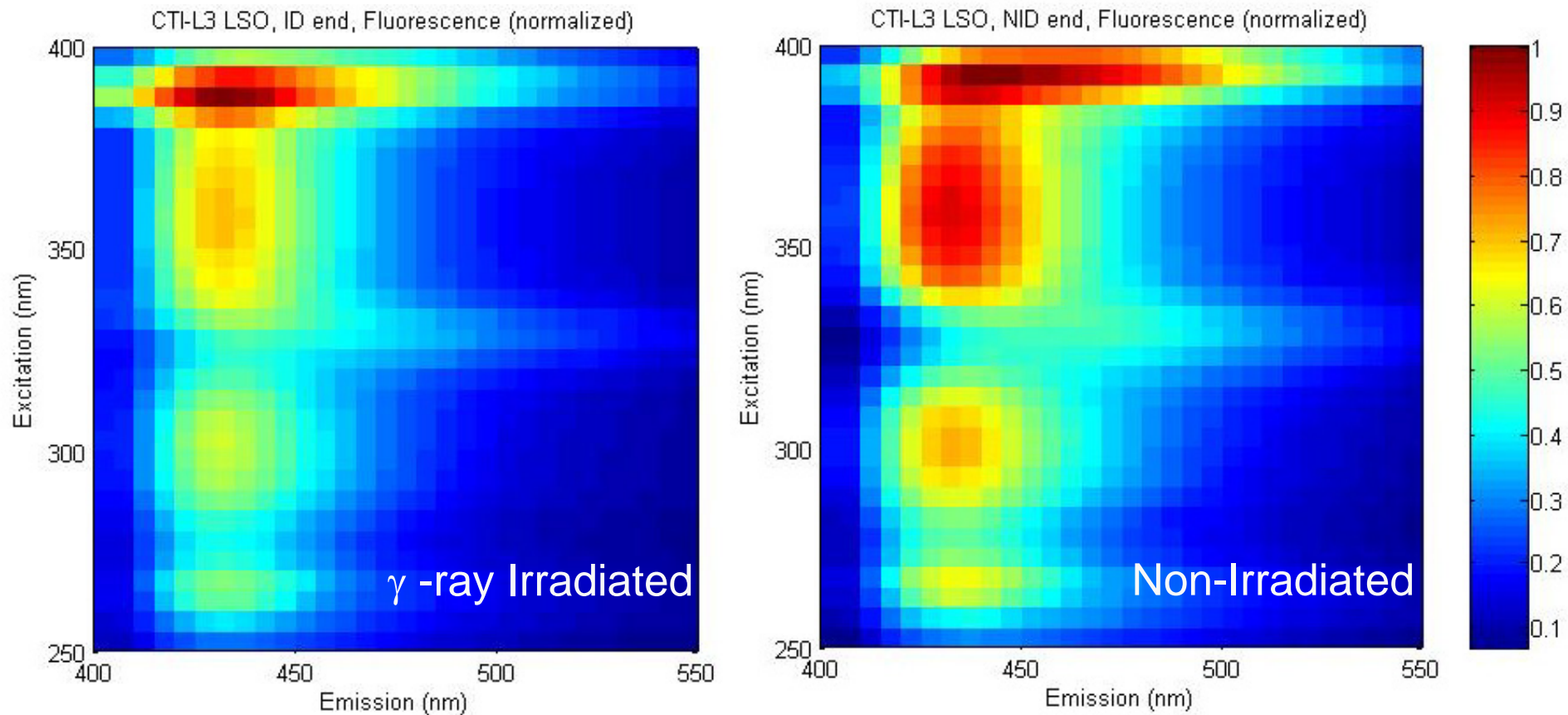
The emission peak of sample's ID (irradiated) end has NO "blue" shift



UV Excited Emission Spectra of Two Halves of the LSO Sample



The γ -ray irradiated half shows less long wavelength emission when excited at 325 nm and 380 nm.

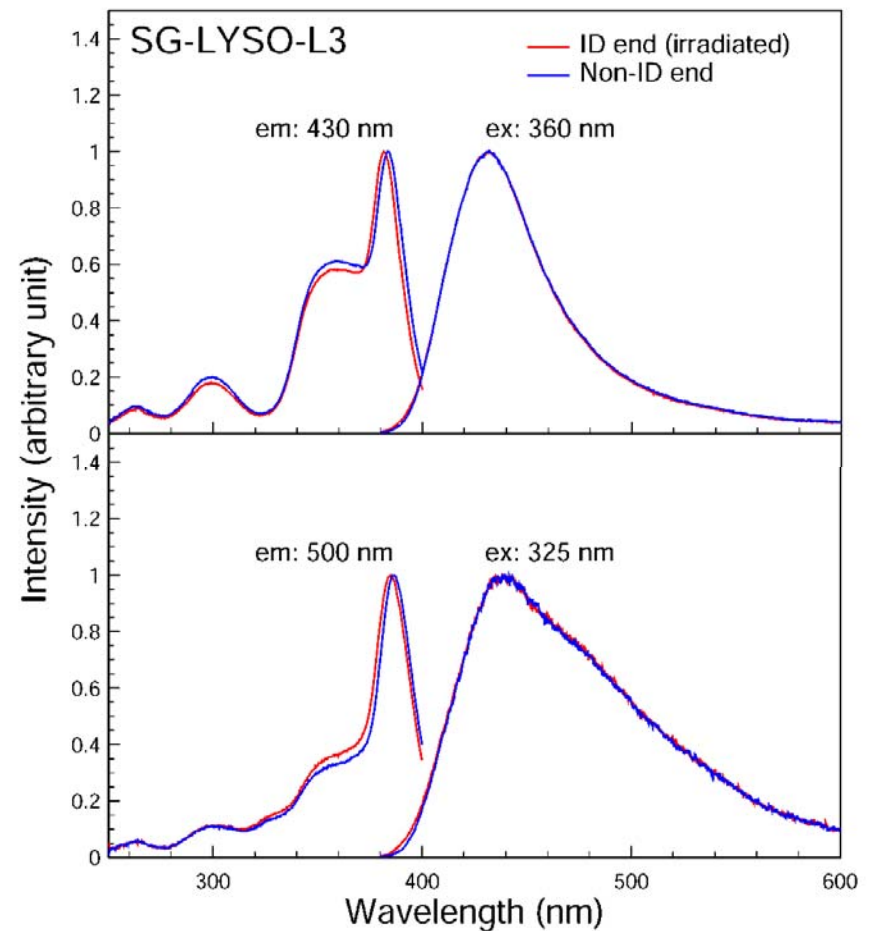
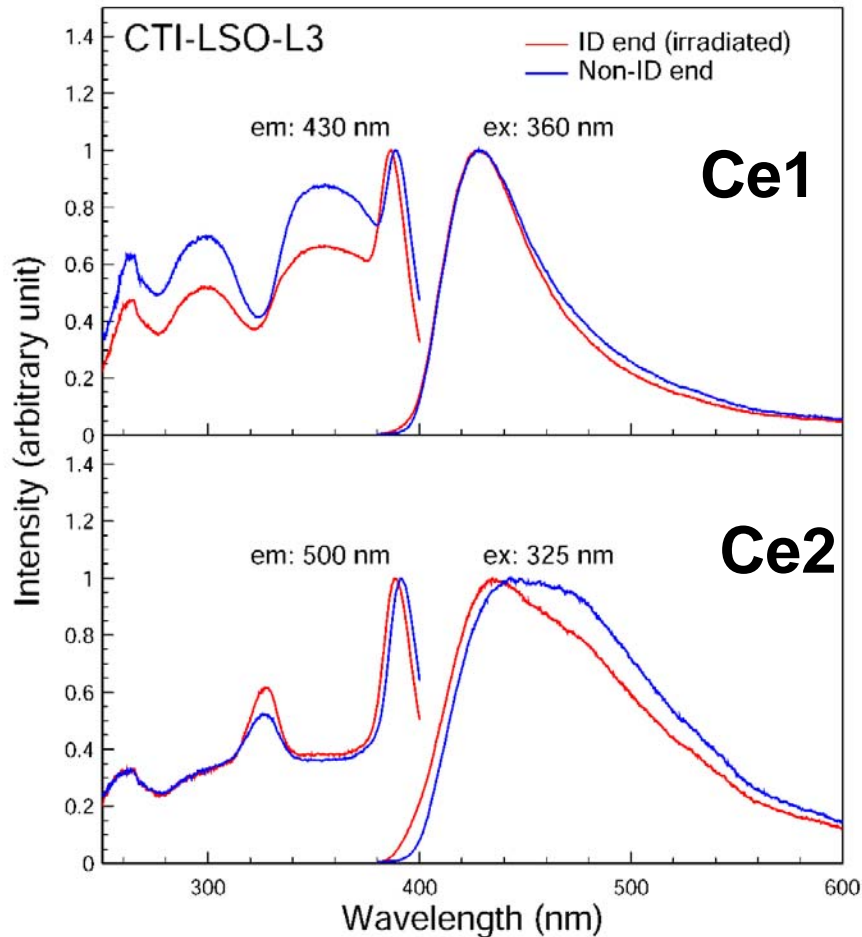


Ce³⁺ Luminescence Centers in LSO



J.D. Naud et. al., IEEE Trans. Nuclear Sci., Vol.43, p1324, June 1996

Ce1: two regular Lu³⁺ crystallographic sites, ex: 360 nm, em: 430 nm
 Ce2: irregular sites, proposed “interstitial site”, ex: 325 nm, em: 500 nm

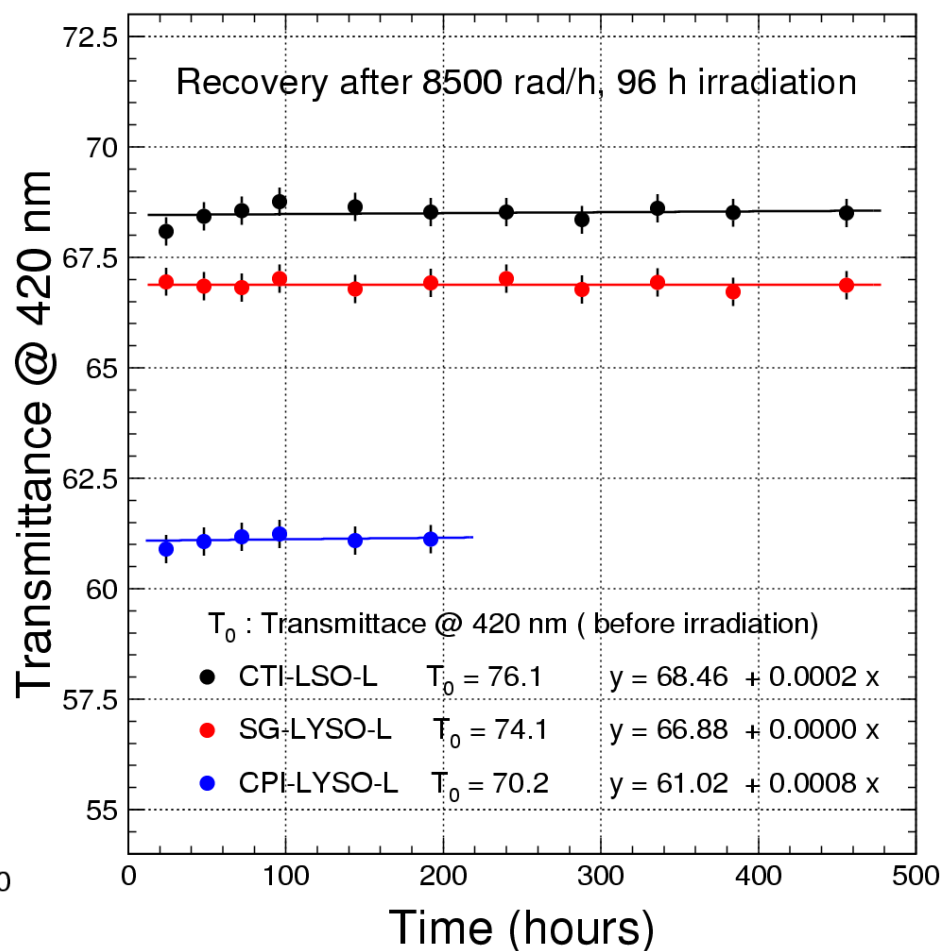
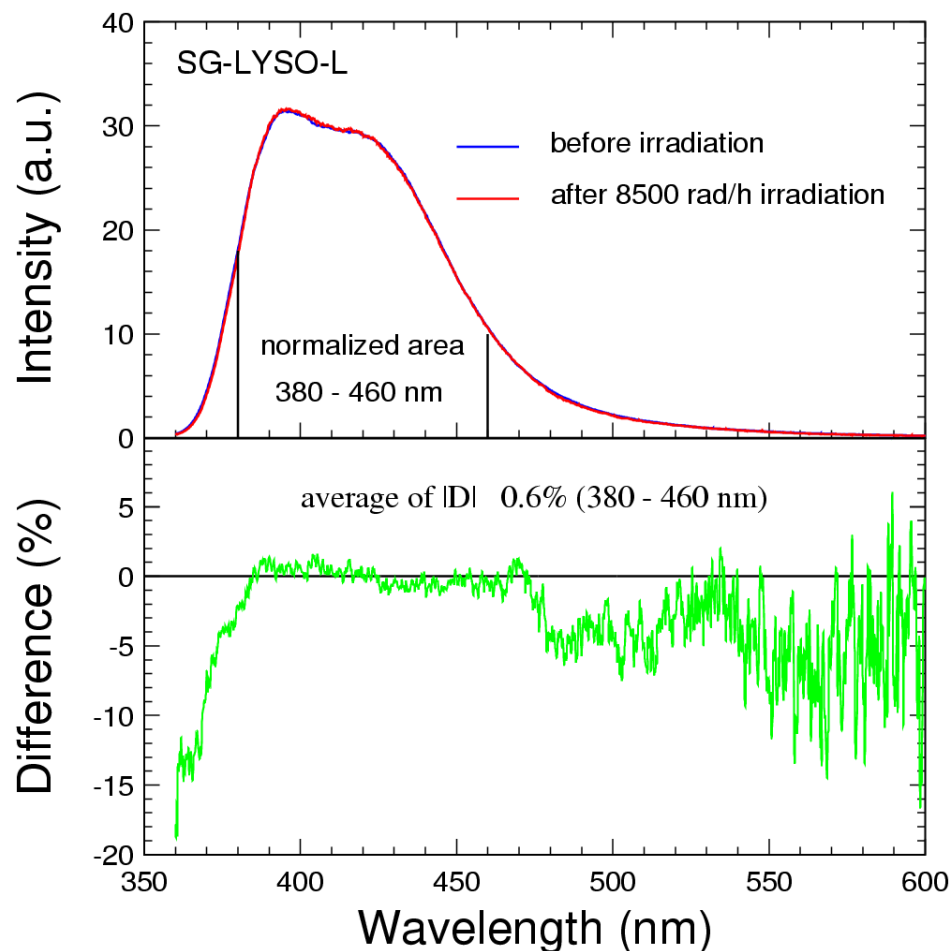


γ -Rays Induced Damage in LSO/LYSO



No damage in Photo-Luminescence

LT recovery very slow

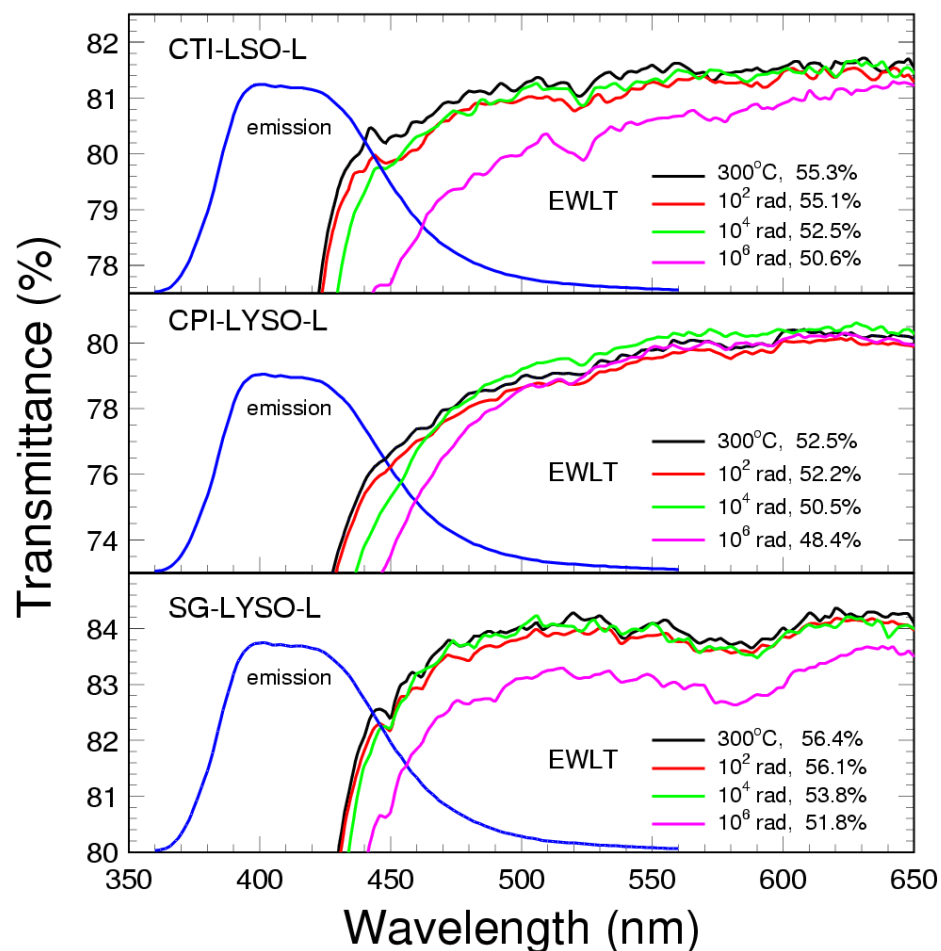
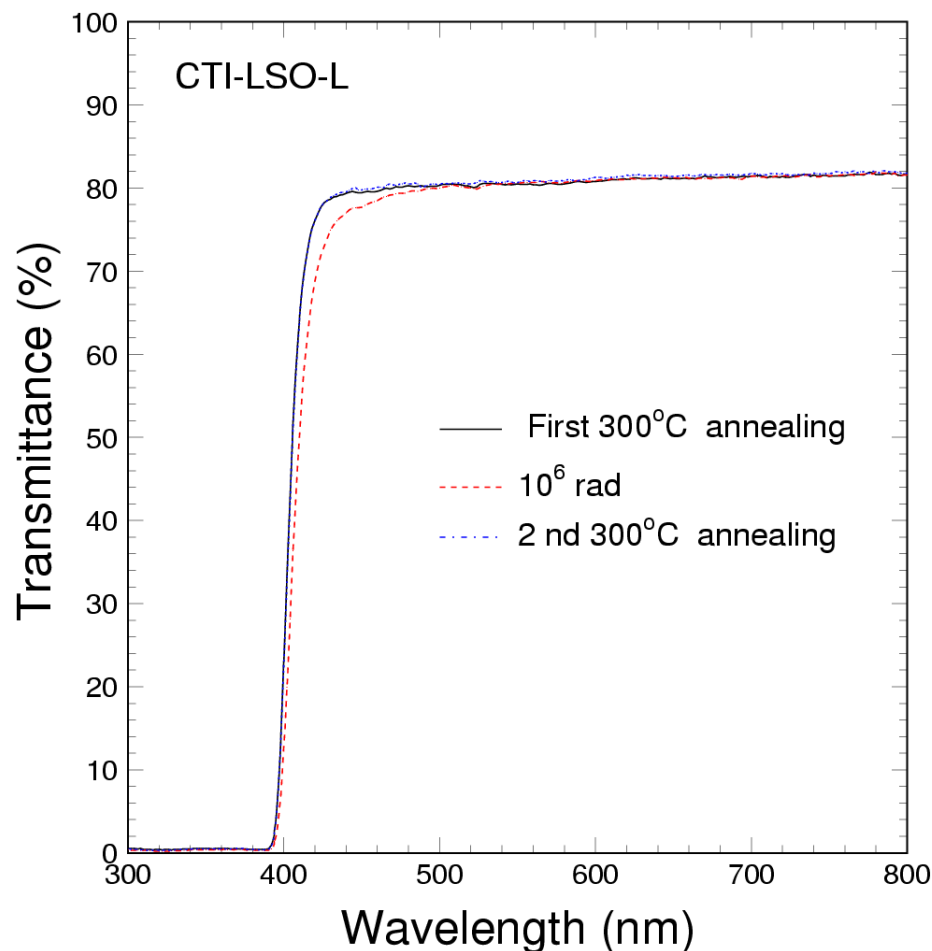


Transmittance Damage



300°C thermal annealing effective

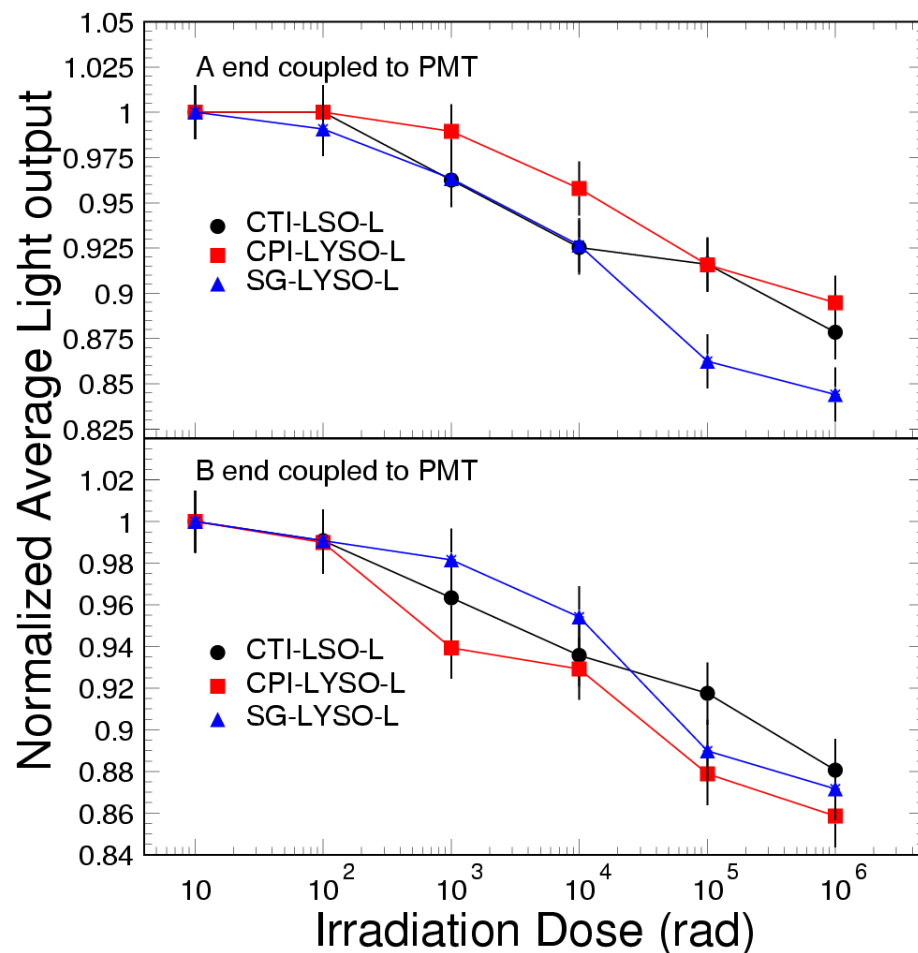
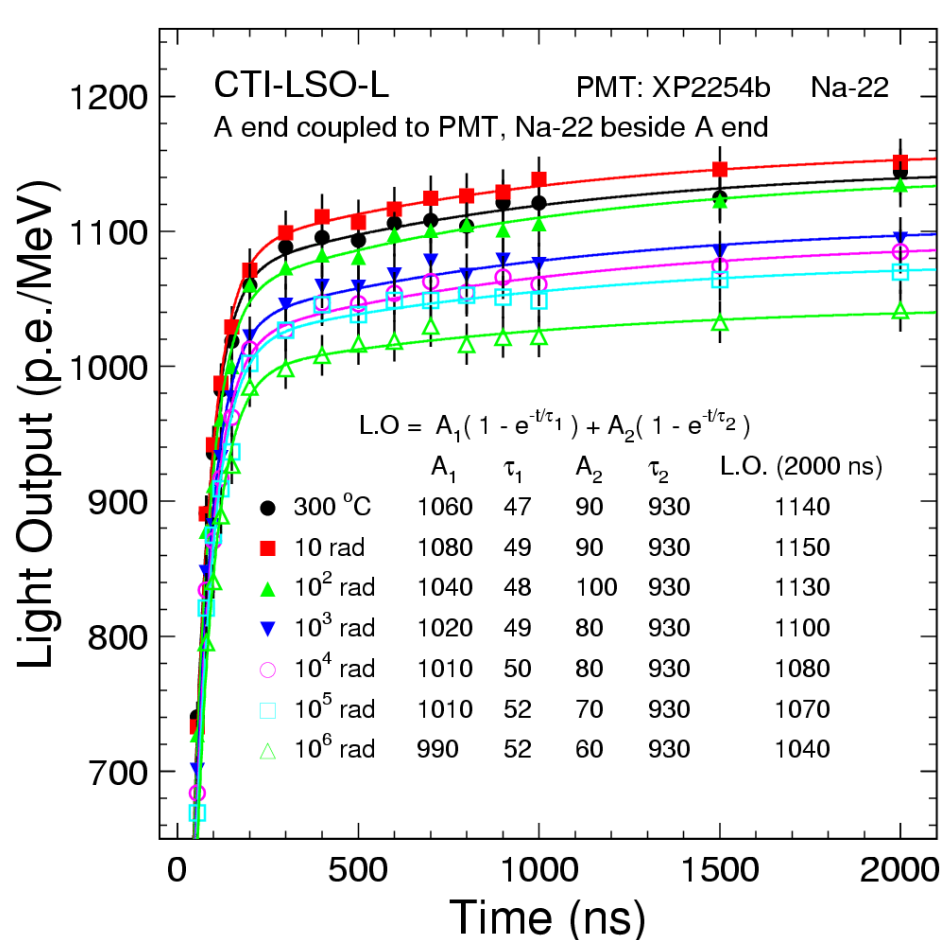
LT damage: 8% @ 1 Mrad



Light Output Damage with PMT



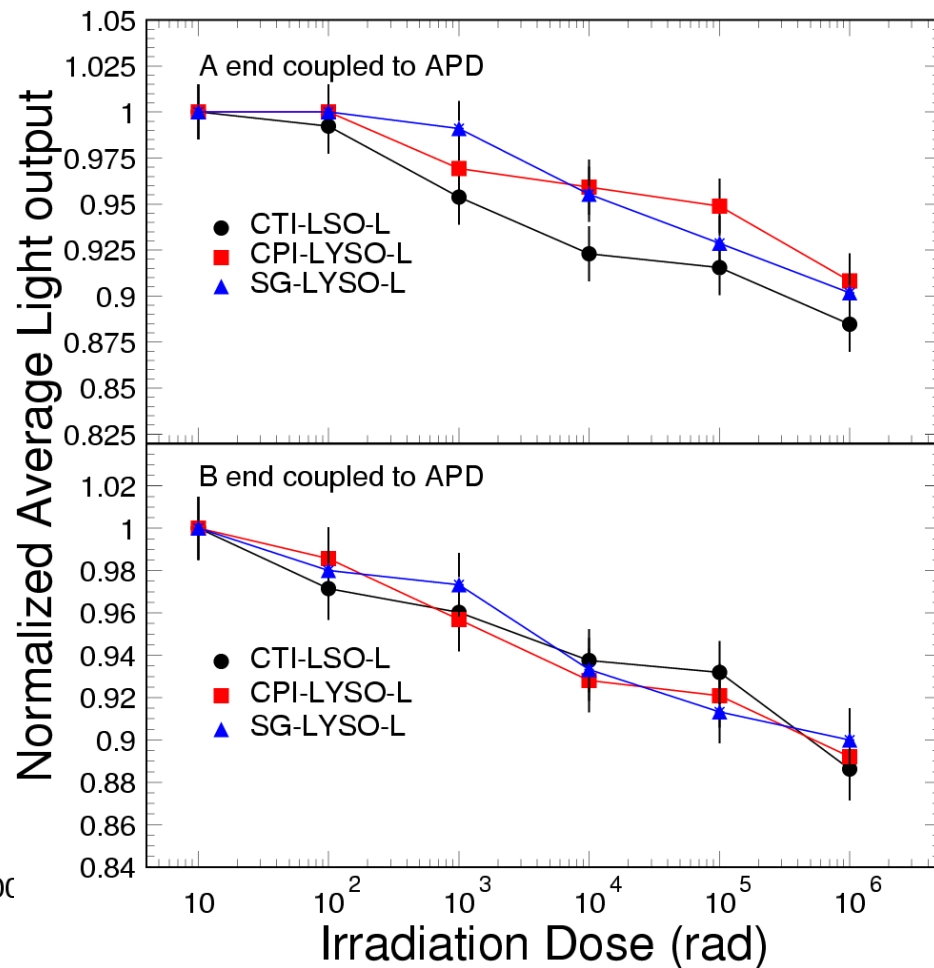
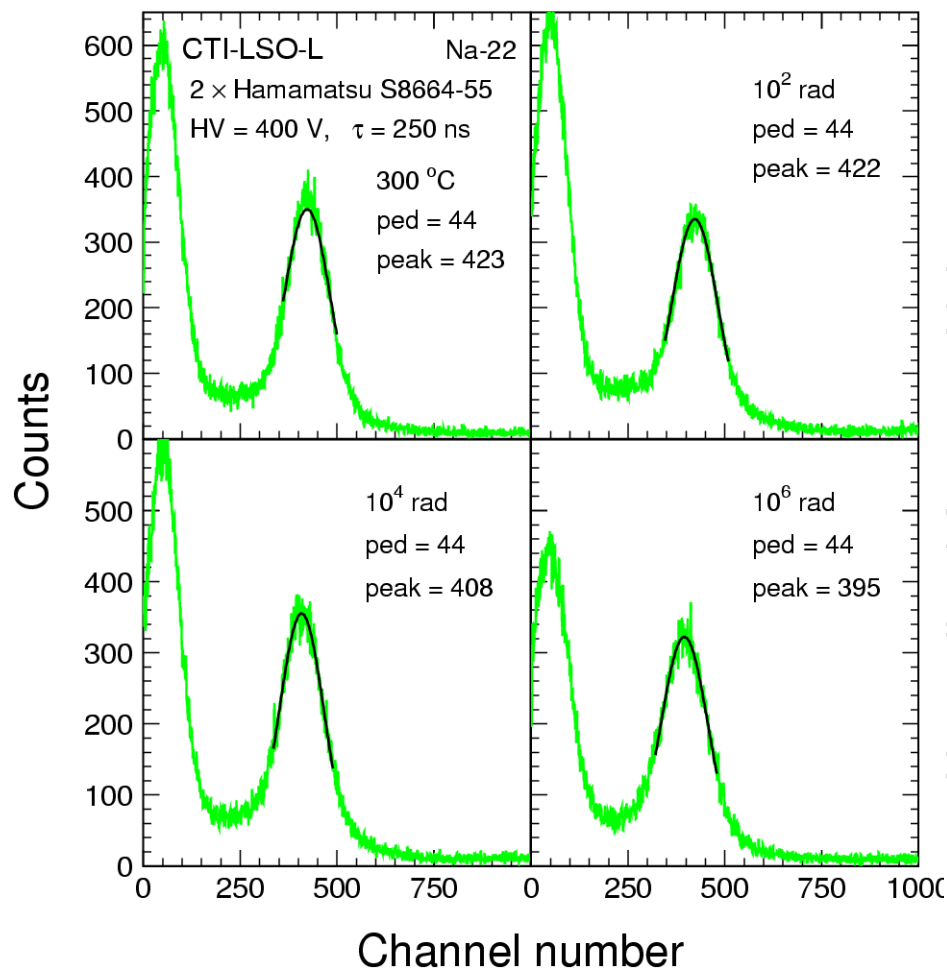
Light output loss: about 12% to 14% @ 1 Mrad



Light Output Damage with APD



Light output loss: about 10% to 12% @ 1 Mrad



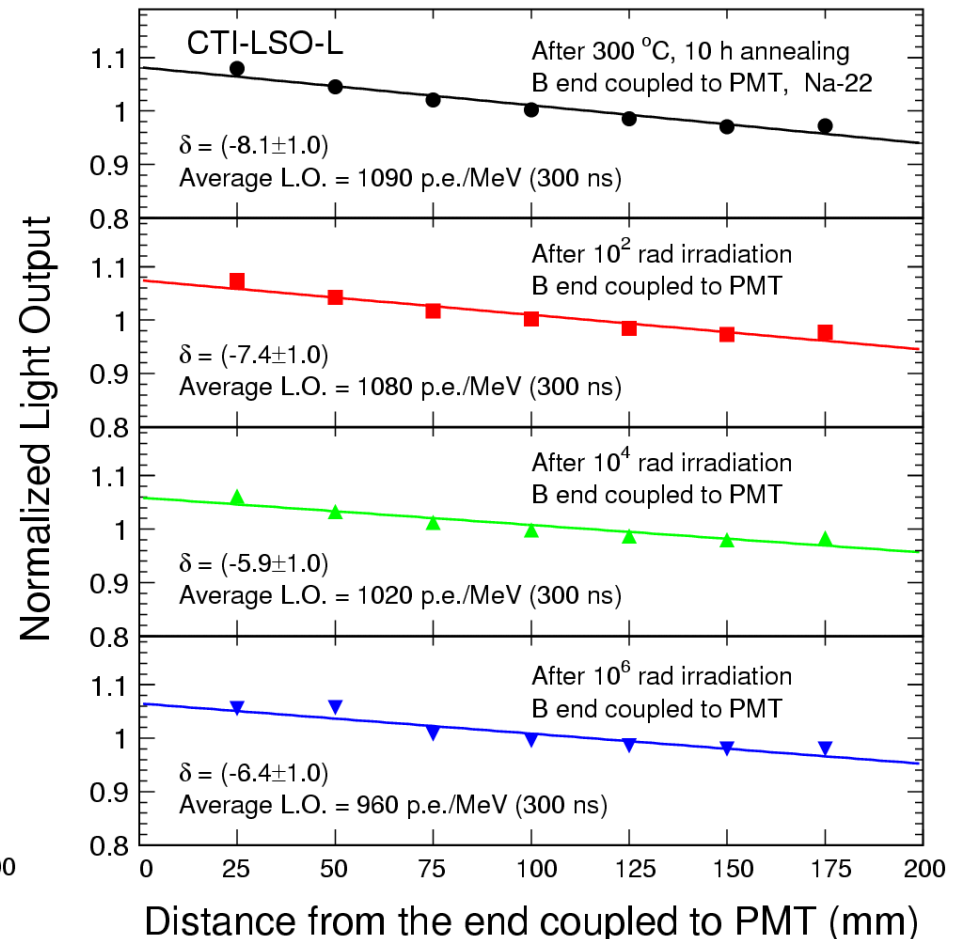
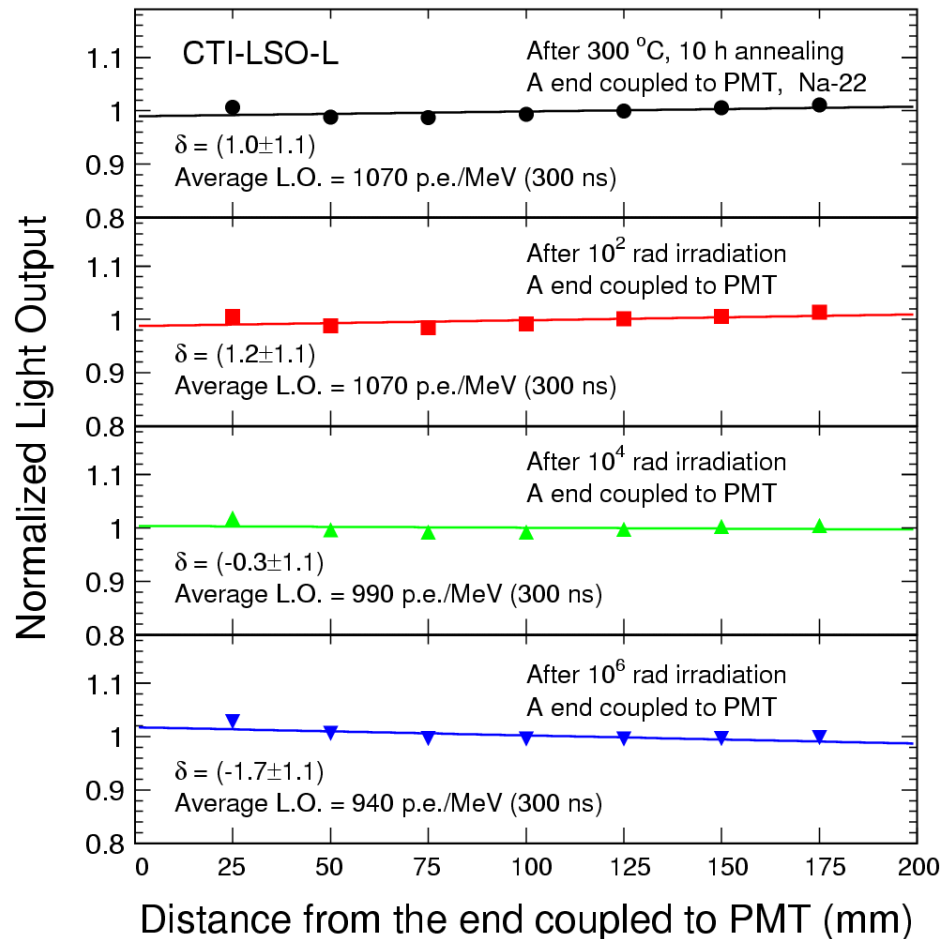
LRU Damage with PMT



Uniformity depends on end coupled to the PMT
No damage in the light response uniformity

A end Coupling

B end Coupling



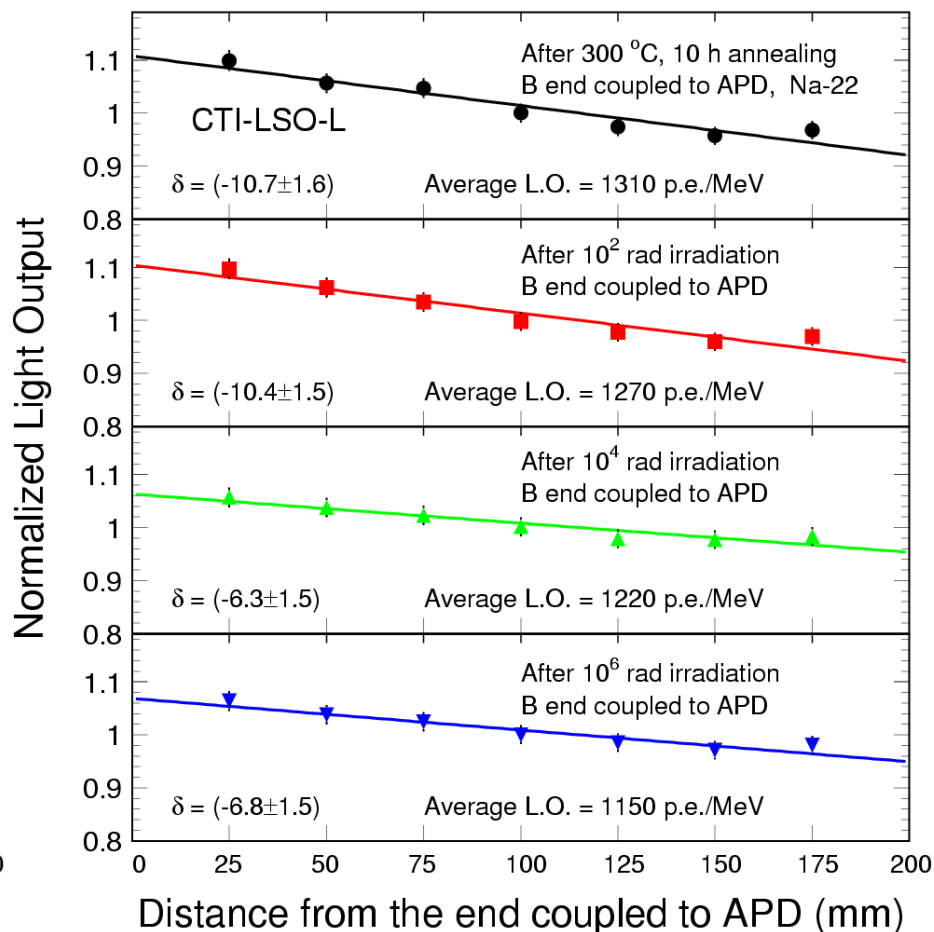
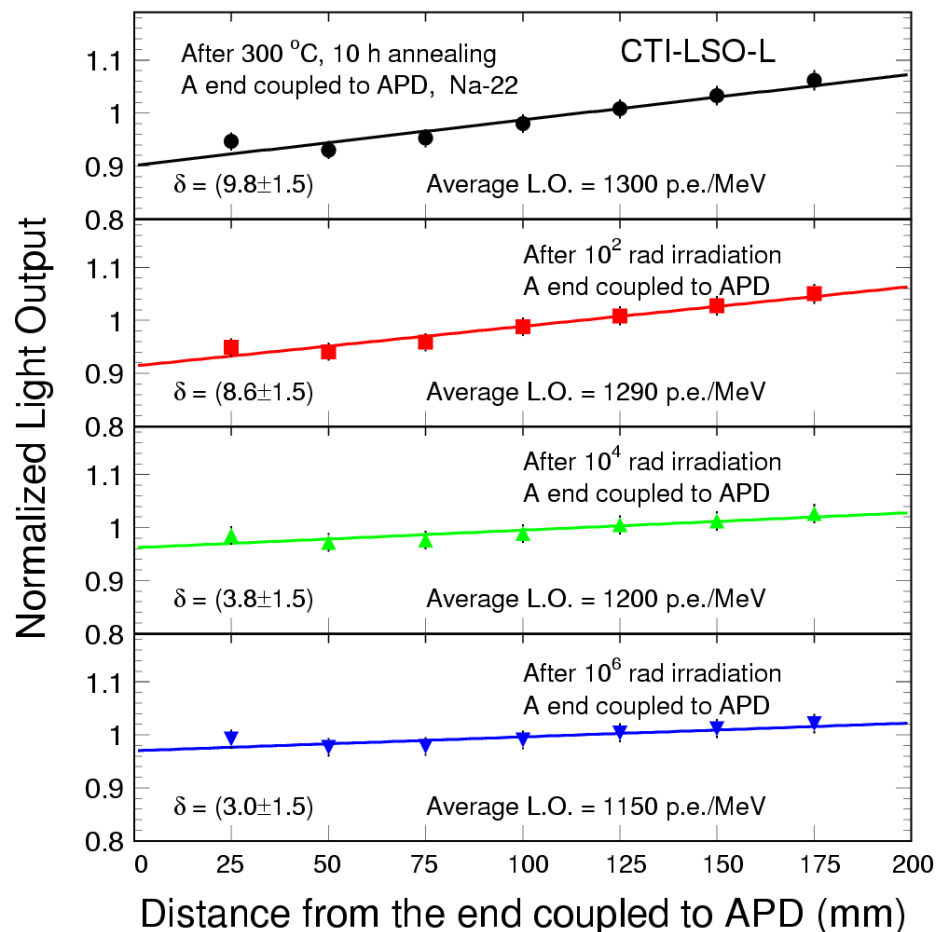
LRU Damage with APD



Uniformity depends on end coupled to the APD
Some change in the light response uniformity

A end Coupling

B end Coupling

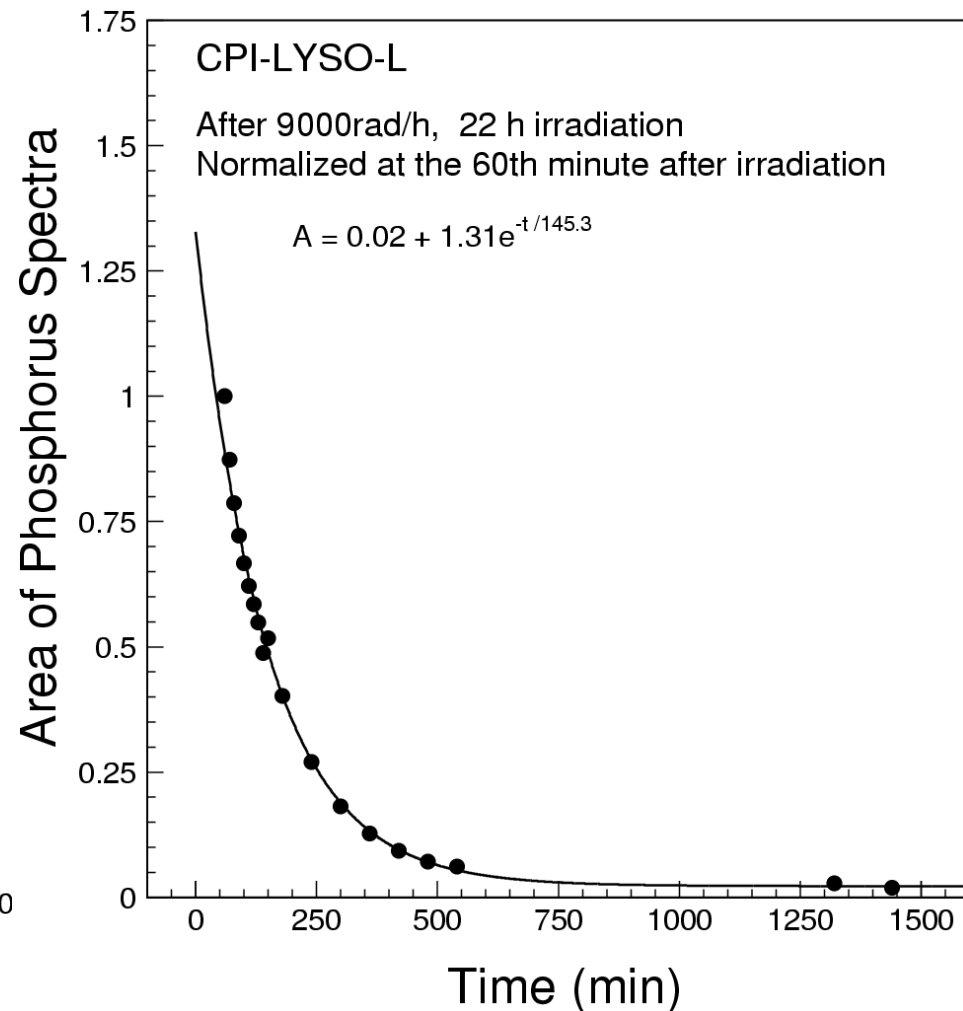
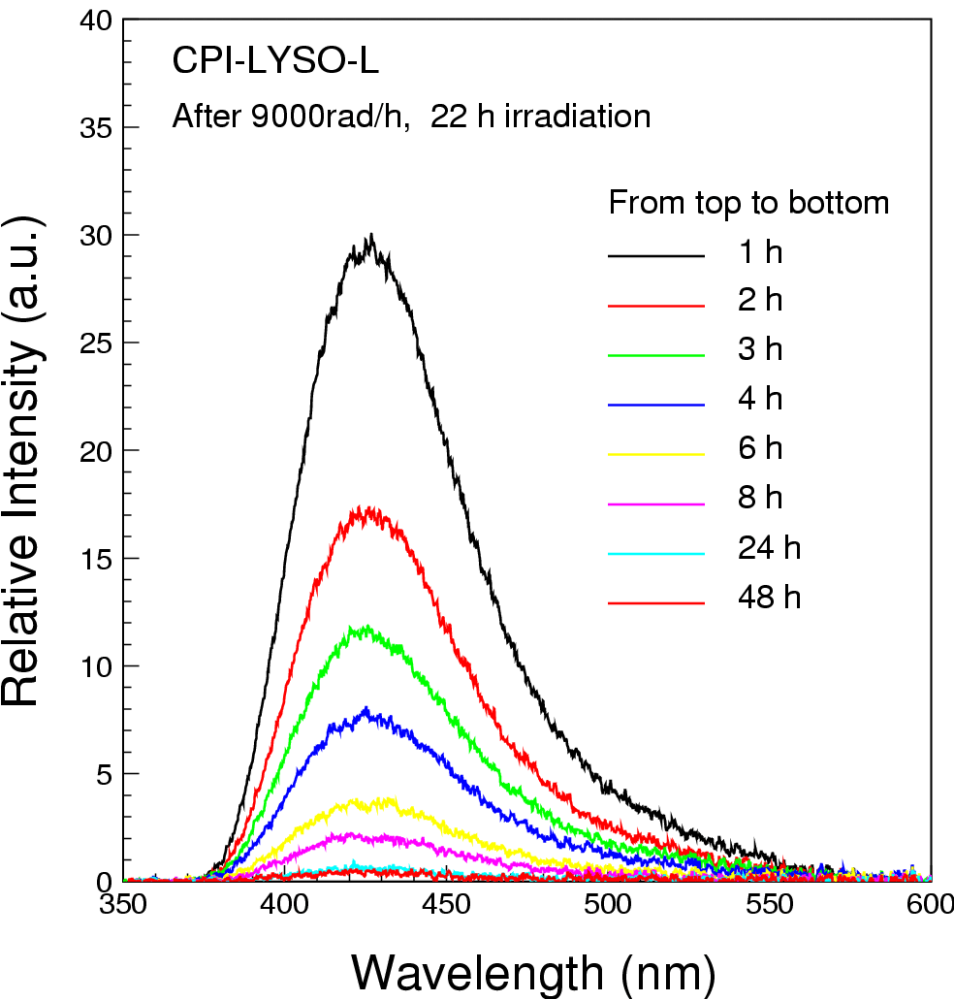




Radiation Induced Phosphorescence



Phosphorescence peaked at 430 nm
with decay time constant of 2.5 h observed

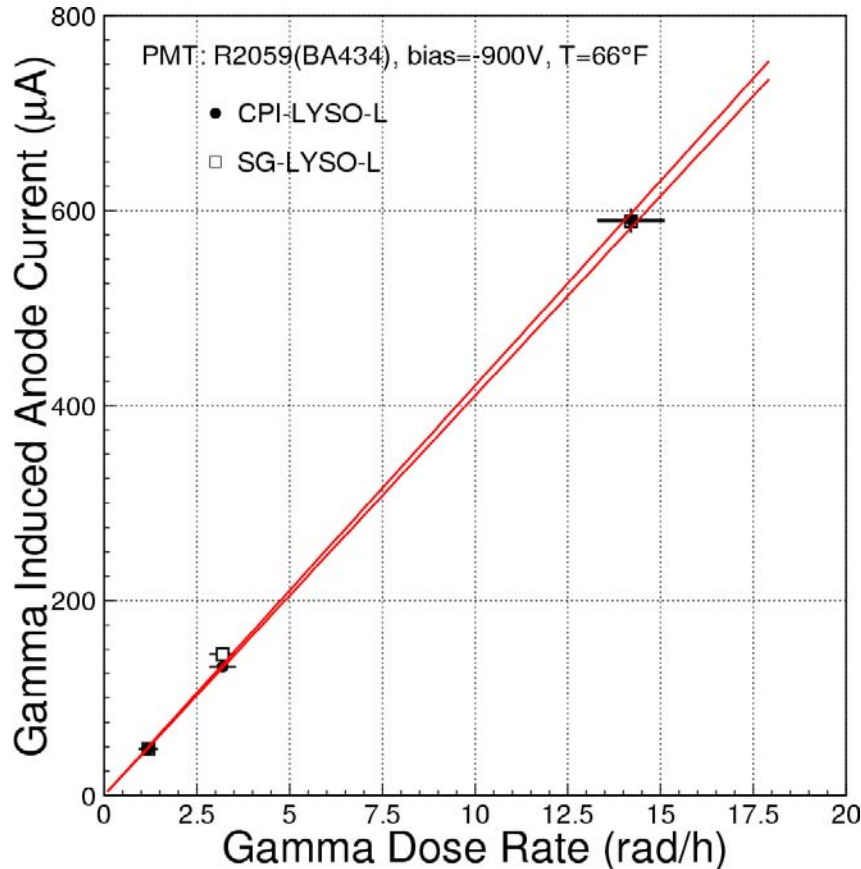




γ -ray Induced Readout Noise



Sample	L.Y.	F	$Q_{15 \text{ rad/h}}$	$Q_{500 \text{ rad/h}}$	$\sigma_{15 \text{ rad/h}}$	$\sigma_{500 \text{ rad/h}}$
ID	p.e./MeV	$\mu \text{ A/rad/h}$	p.e.	p.e.	MeV	MeV
CPI	1,480	41	6.98×10^4	2.33×10^6	0.18	1.03
SG	1,580	42	7.15×10^4	2.38×10^6	0.17	0.97



γ -ray induced PMT anode current can be converted to the photoelectron numbers (Q) integrated in 100 ns gate. Its statistical fluctuation contributes to the readout noise (σ).



LSO/LYSO ECAL Performance



- Less demanding to the environment because of small temperature coefficient.
- Radiation damage is less an issue as compared to other crystals.
- A better energy resolution, $\sigma(E)/E$, at low energies than L3 BGO and CMS PWO because of its high light output and low readout noise:

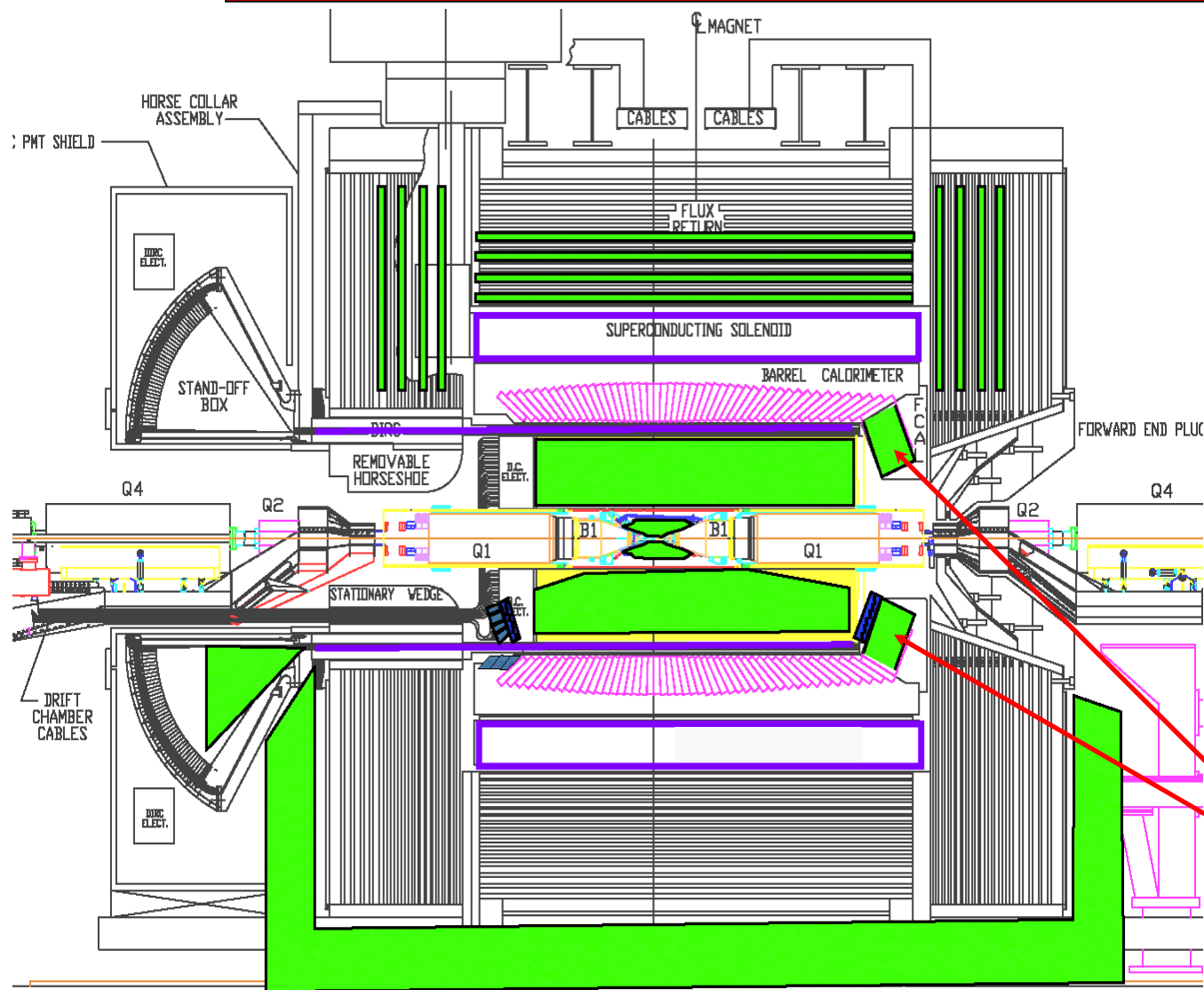
$$2.0\% / \sqrt{E} \oplus 0.5\% \oplus .001/E$$



LYSO Development for SuperB



David Hitlin The SuperB Project N01-3 IEEE NSS 2007



Aiming at $10^{36}/\text{cm}^2/\text{s}$ luminosity for rare B decays

Need fast detector with low noise at endcap

LSO/LYSO



LSO/LYSO Mass Production



CTI: LSO



CPI: LYSO



Saint-Gobain
LYSO



Additional Capability: SIPAT @ Sichuan, China



Six LSO & LYSO Samples



2.5 x 2.5 x 20 cm (18 X_0) Bar



Three CTI LSO samples are provided by Chuck Melcher.

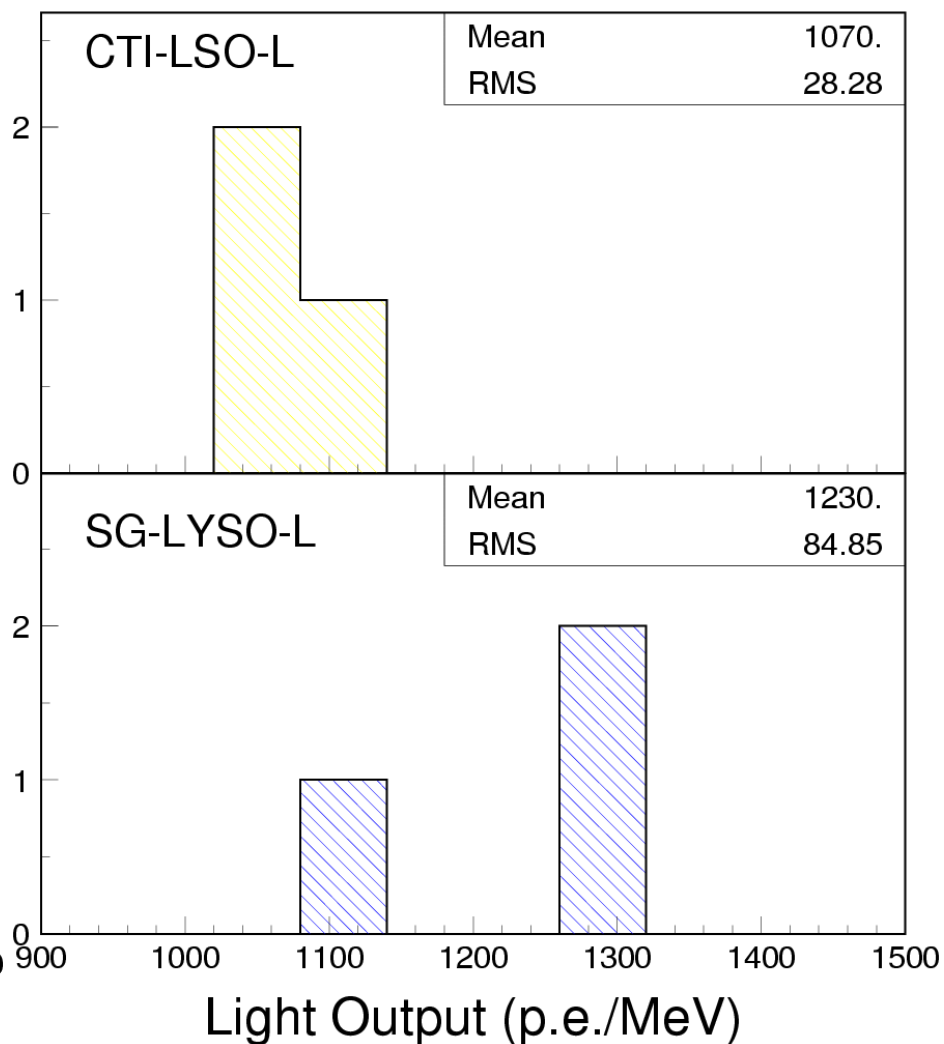
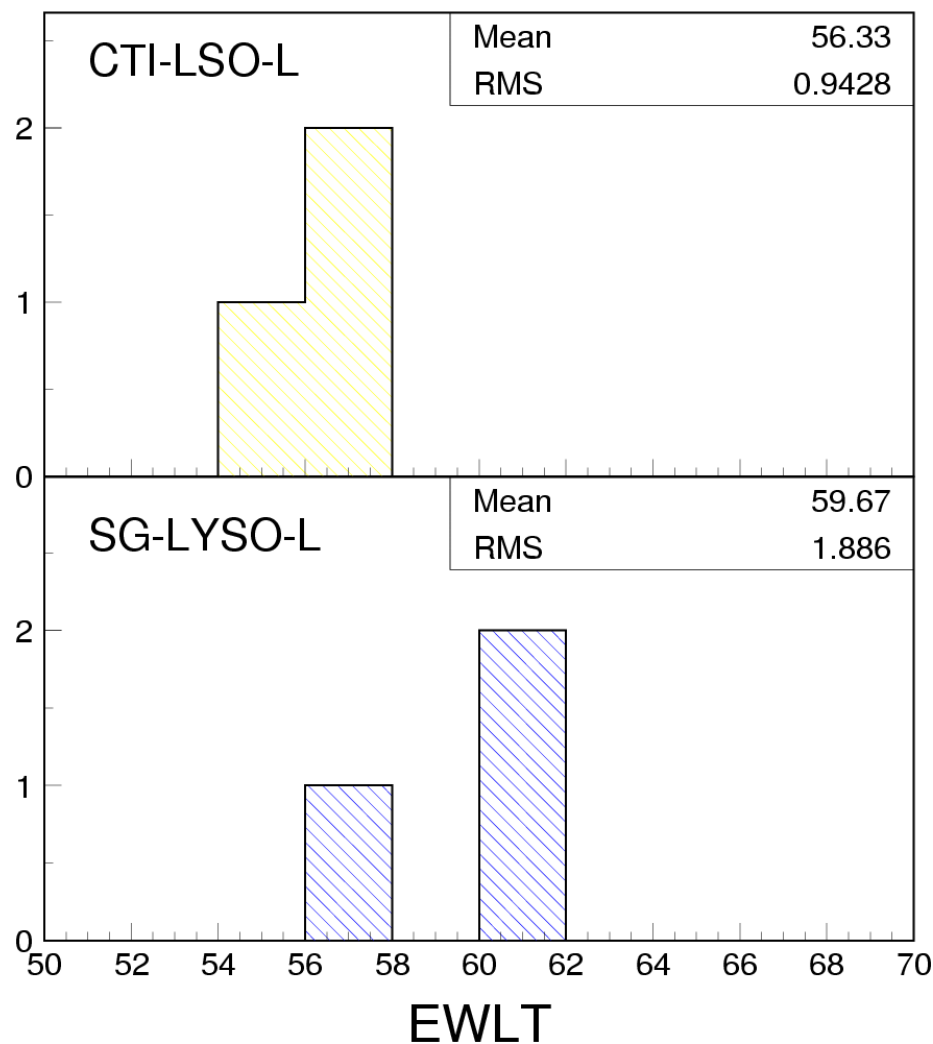
Three LYSO samples are purchased from Saint-Gobain.



Statistical Comparison



Recent LYSO crystals are better than LSO





Sichuan Institute of Piezoelectric and Acousto-optic Technology (SIPAT)



China Electronics Technology Corporation (CETC)
No. 26 Research Institute, www.sipat.com



SIPAT R&D Building



Total area: 52,258m²

Construction area: 38,331m²

For R&D : 27,765m²



SIPAT: Furnace & R&D Issues



- Raw material:
 Lu_2O_3 : 99.995%
 SO_2 : 99.999%
- Stoichiometry
- Temperature Gradient
- Growth Parameter Optimization
- Thermal Annealing
- Iridium Crucible Maintenance
- Power Supply Stability
- Chilled Water Stability



LYSO Growth Progress at SIPAT



Started 2001, invested >\$1M, Significant Progress in last year

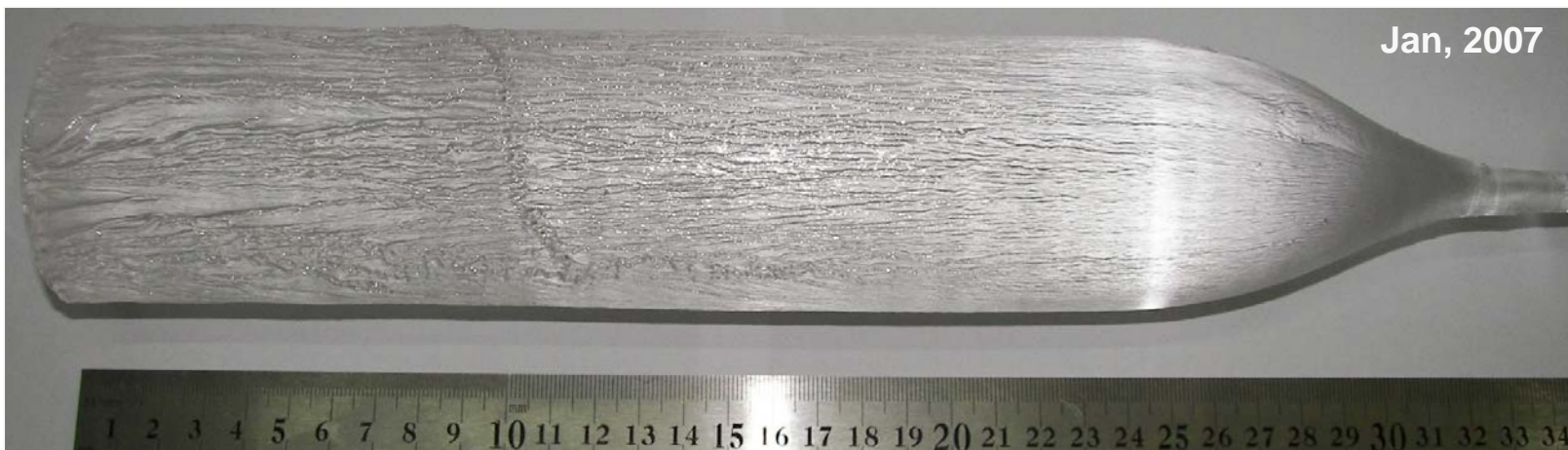
May, 2005



Sep, 2006



Jan, 2007





SIPAT \varnothing 60 x 250 mm LYSO Boles

Sep, 2007





SIPAT Czochralski Furnaces





First SIPAT LYSO Sample for HEP



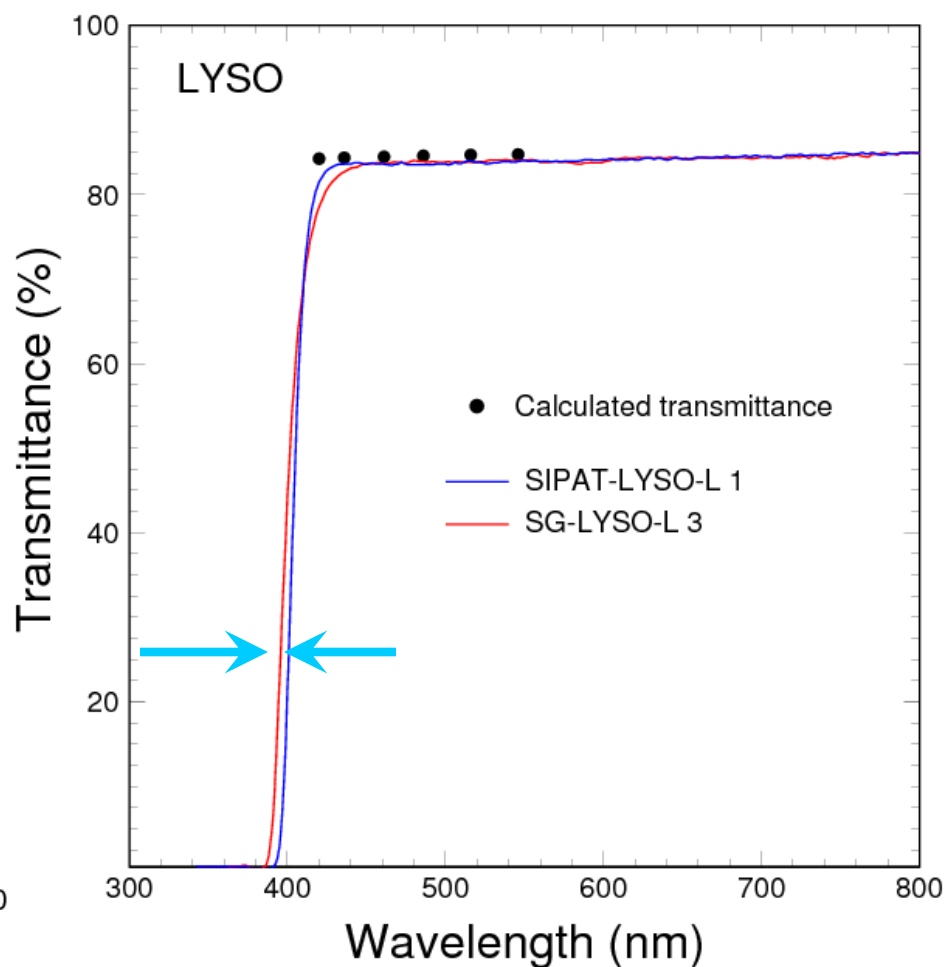
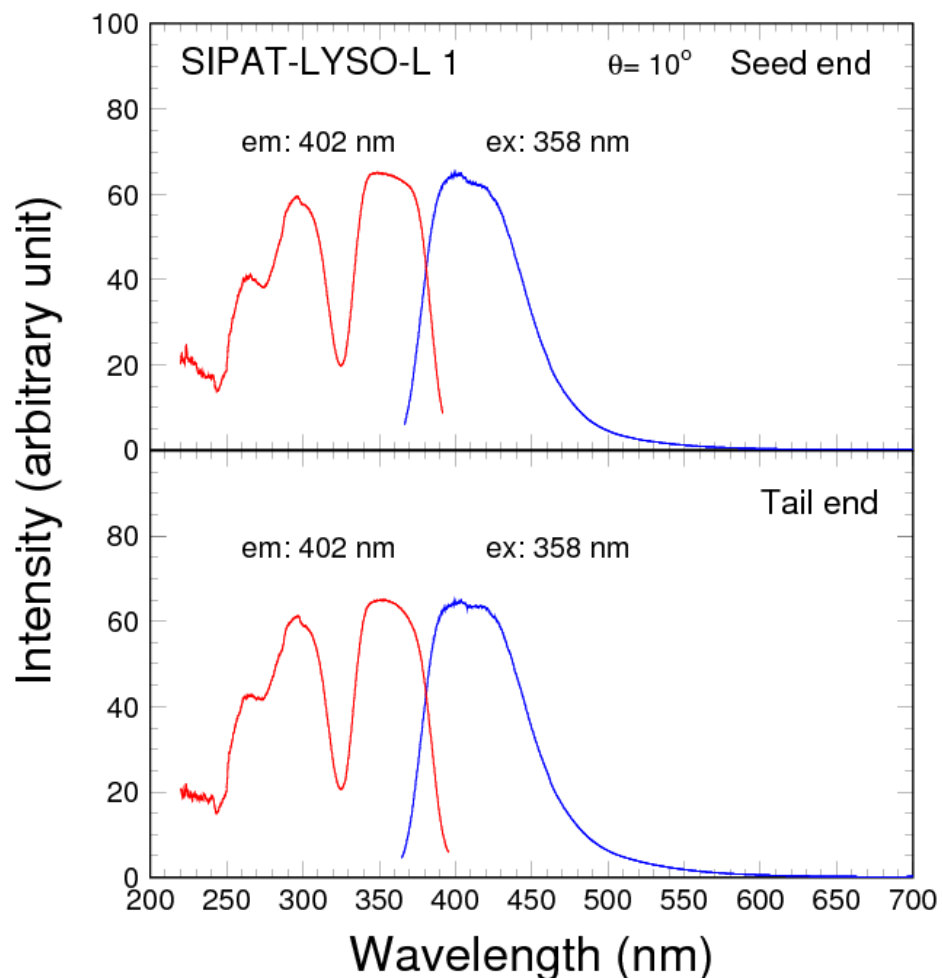
- Received in the middle of August with dimension of 25 x 25 x 200 mm and good visual inspection.
- It was first annealed at 300°C for 10 hours and with its initial optical and scintillation properties measured.
- Together with SG-L3, two samples were irradiated with integrated doses of 10, 10^2 , 10^3 , 10^4 , 10^5 and 10^6 .
- Samples were kept in dark after irradiation for 48 hours before optical and scintillation property measurement.
- Damage to transmittance, light output and uniformity are compared with samples from CTI, CPI and Saint-Gobain.

Initial Optical Properties



Excitation: emission @ 402 nm
Emission: excitation @ 358 nm

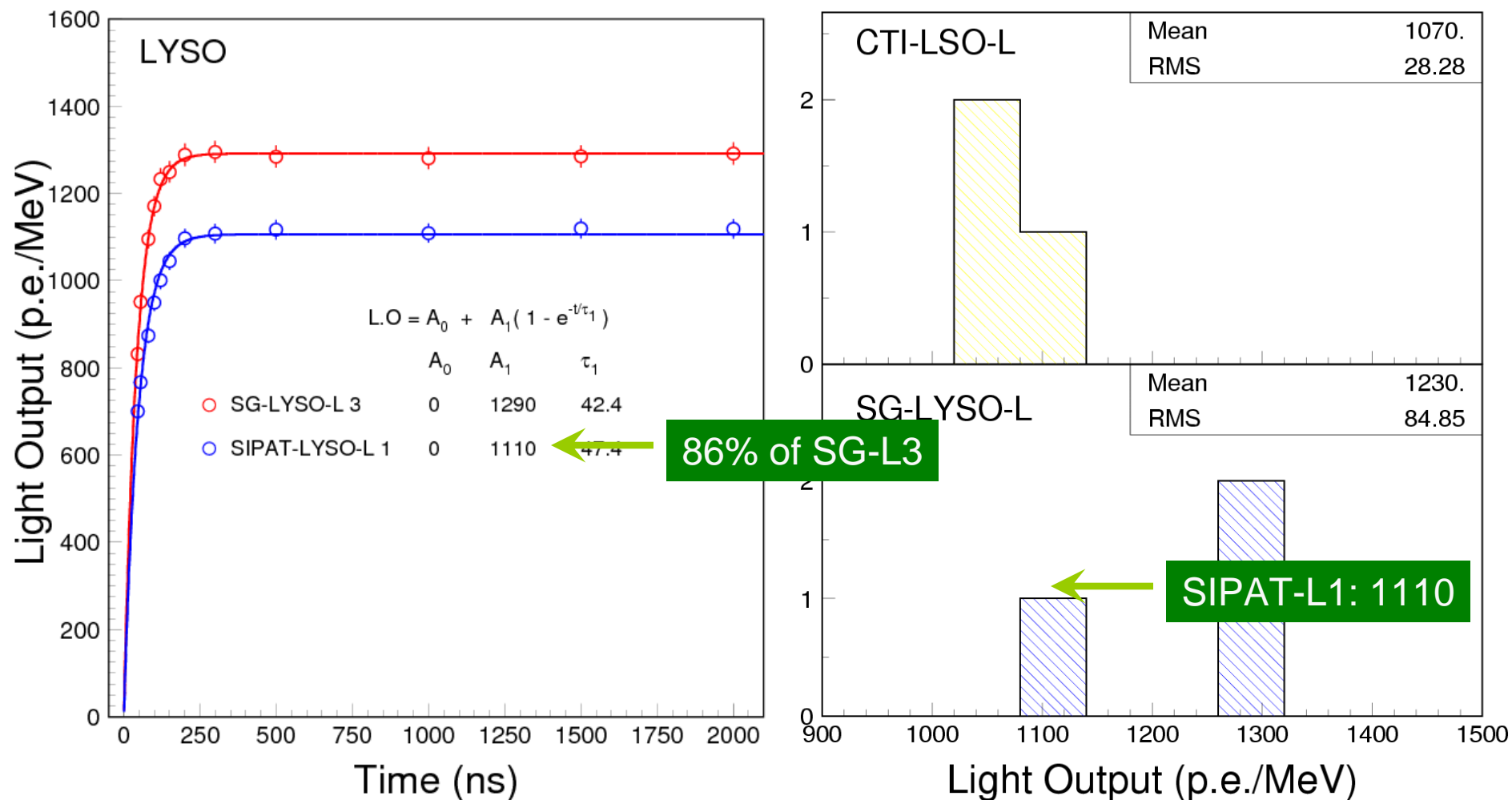
The cutoff of SG-L3 has ~5 nm blue shift compared to SIAPT-L1



Light Output & Decay Kinetics



Compatible with the first batch large size samples from CTI and Saint-Gobain, and is 86% of the 'best' samples



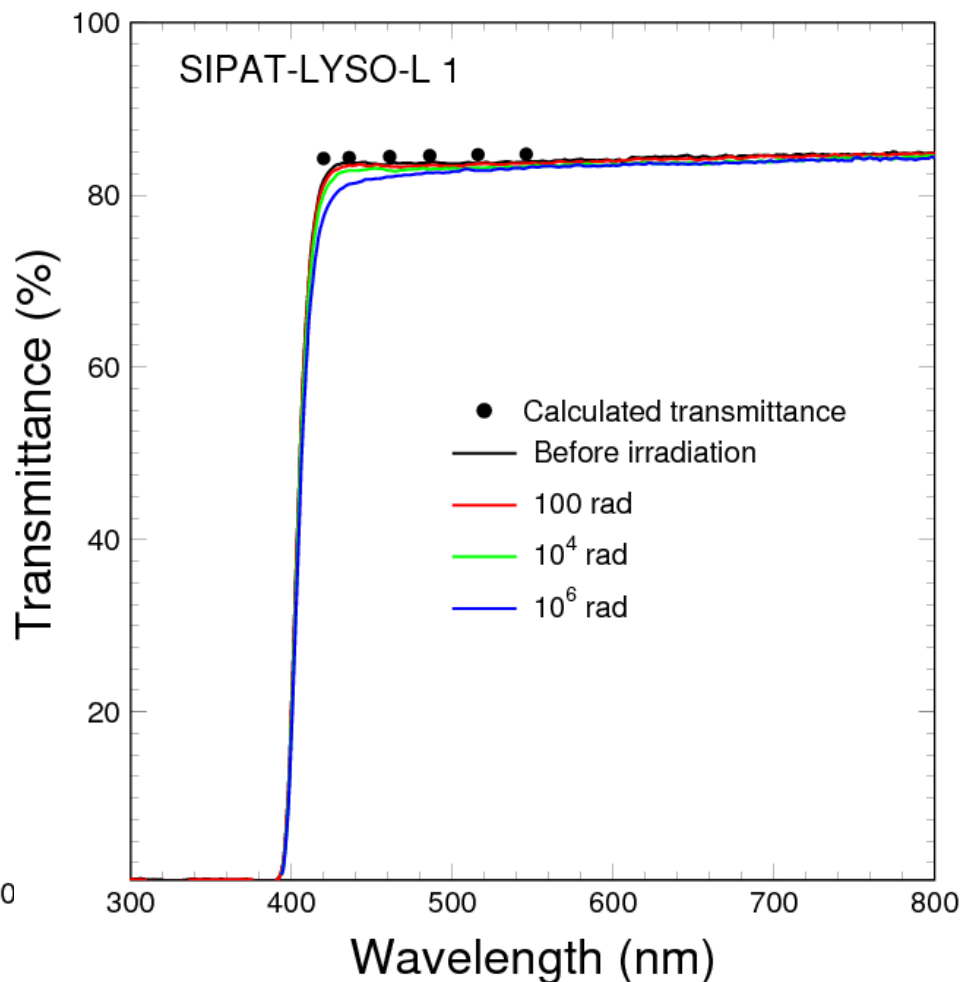
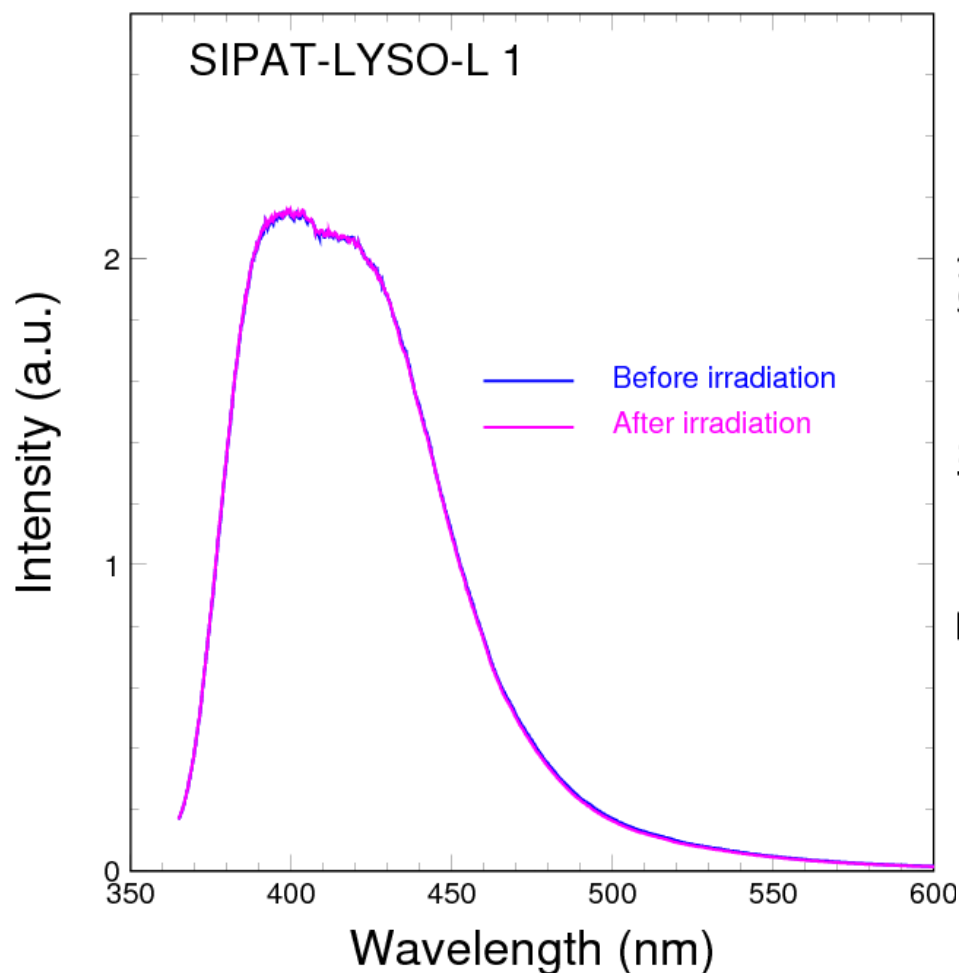


γ -Ray Induced Radiation Damage



Scintillation spectrum
not affected by irradiation

~8% damage @ 420 nm
after 10^6 rad irradiation



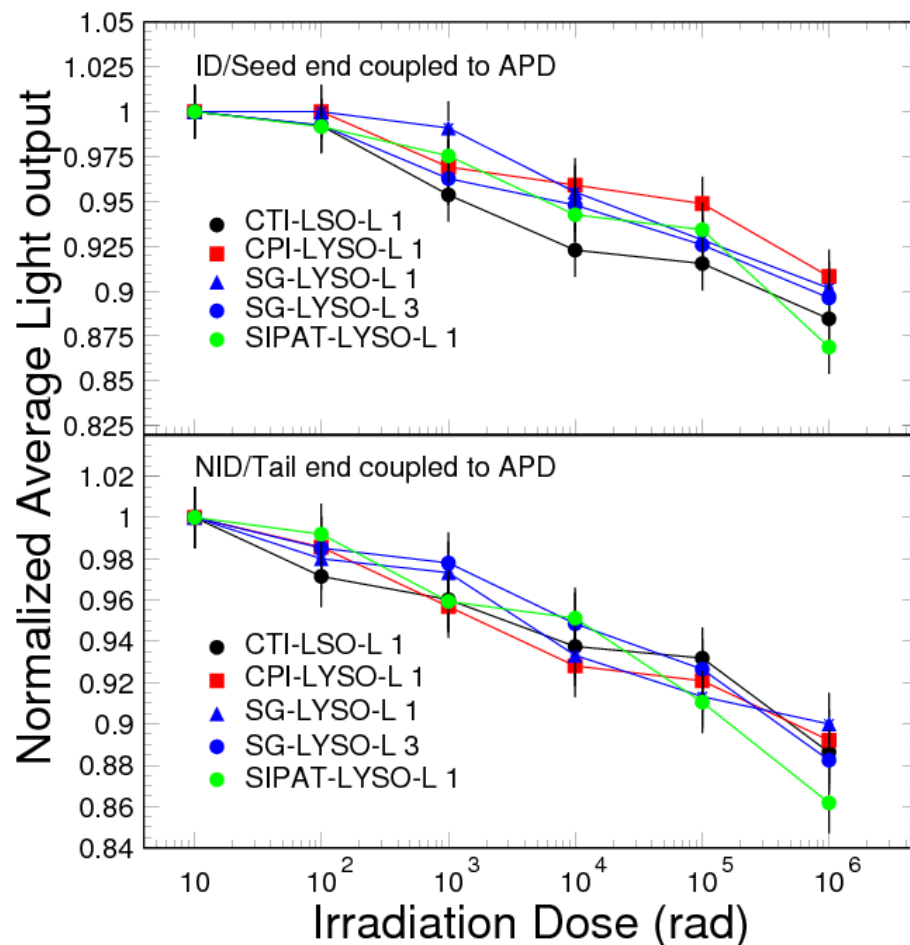
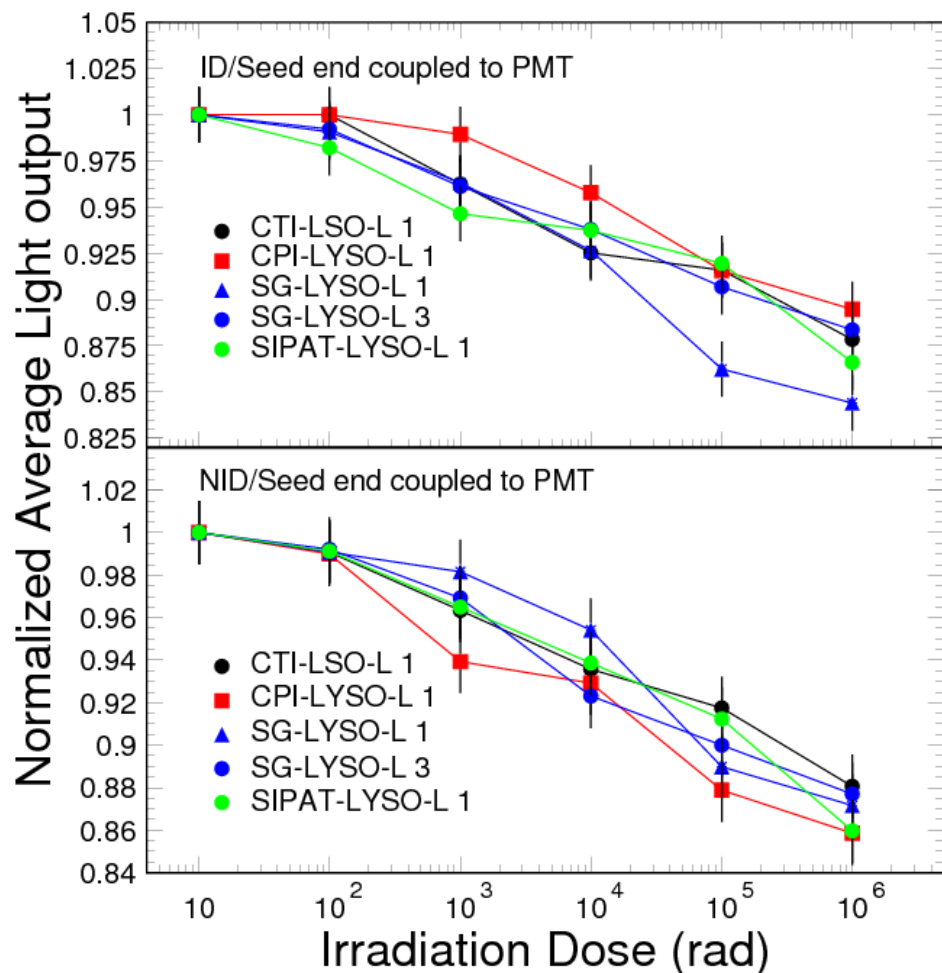
Comparison of L.O. Damage



All samples show consistent radiation resistance

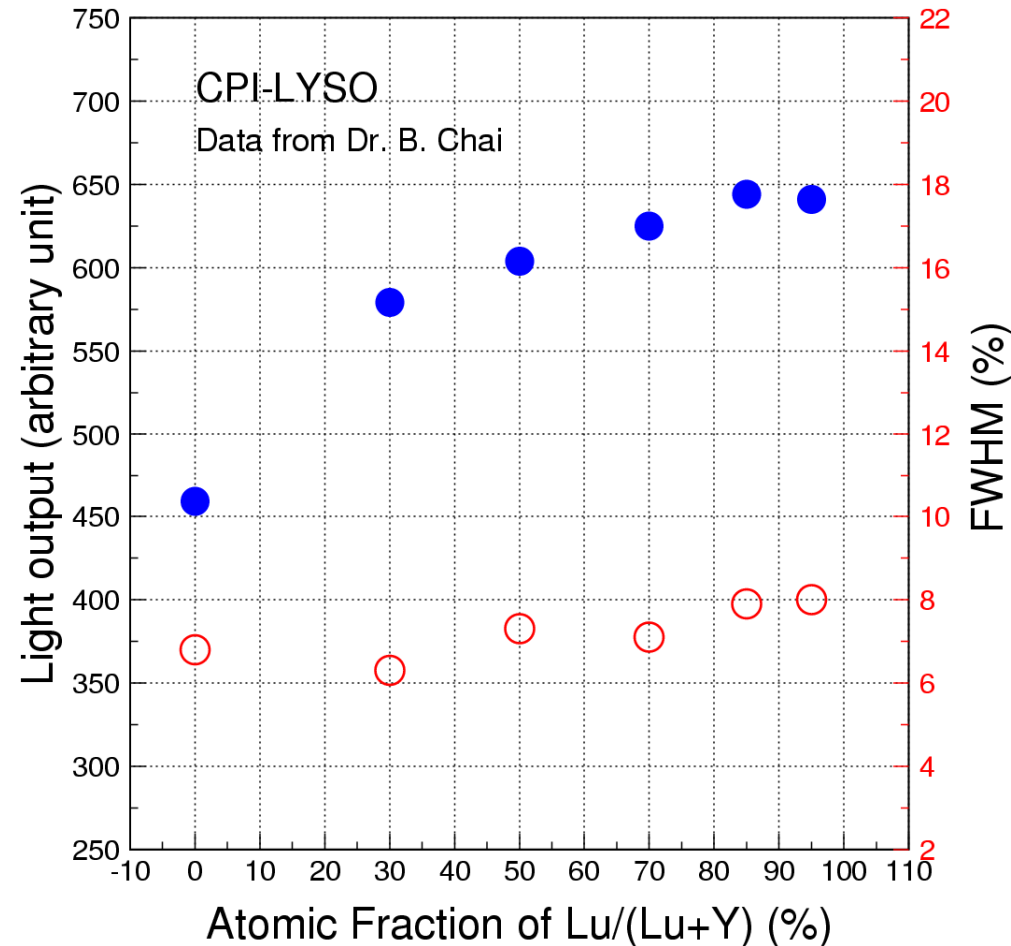
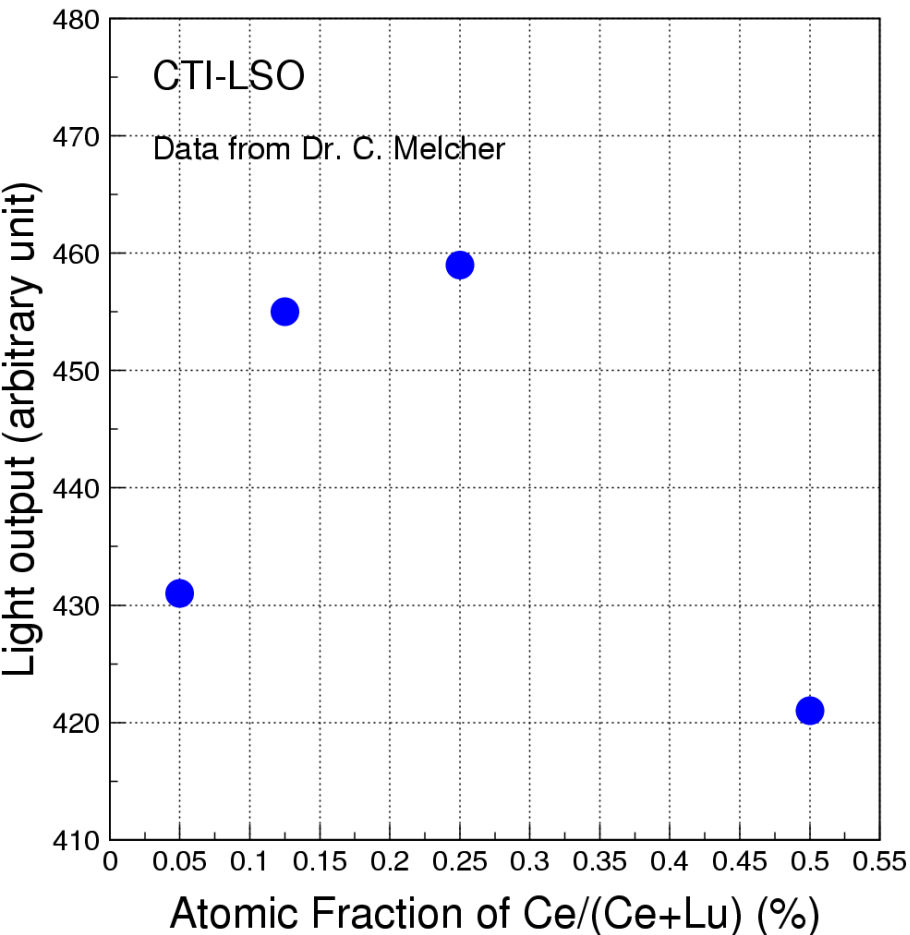
10% - 15% loss by PMT

9% - 14% loss by APD



Possible Origin of Non Uniformity

C. Melcher: LO in LSO is a function of Ce concentration
B. Chai: LO in LYSO is a function of Yttrium fraction

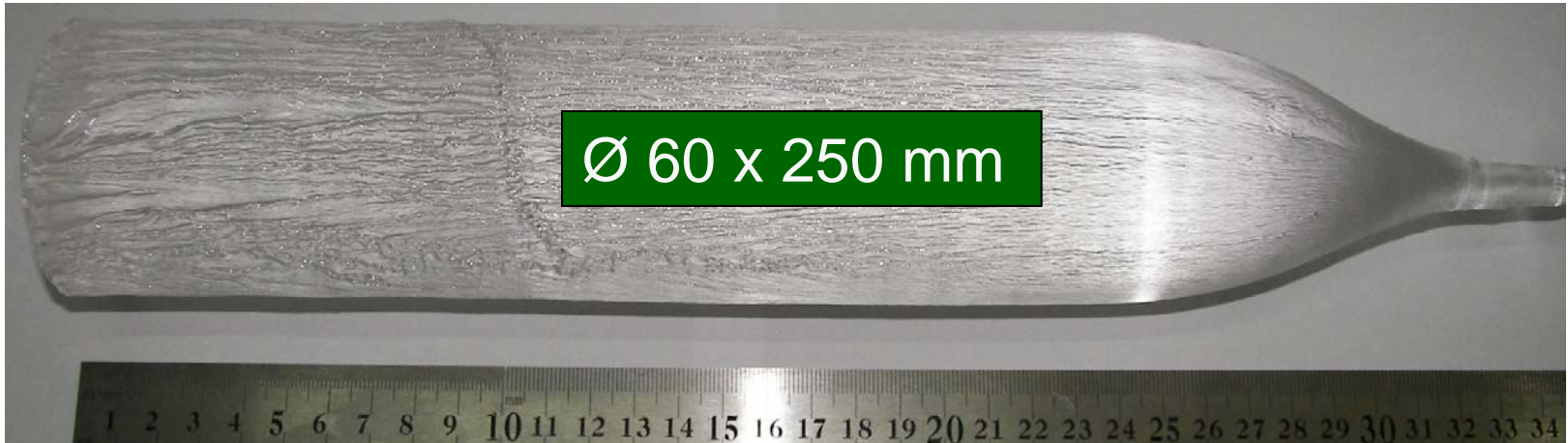




First SIPAT LYSO Bole for HEP



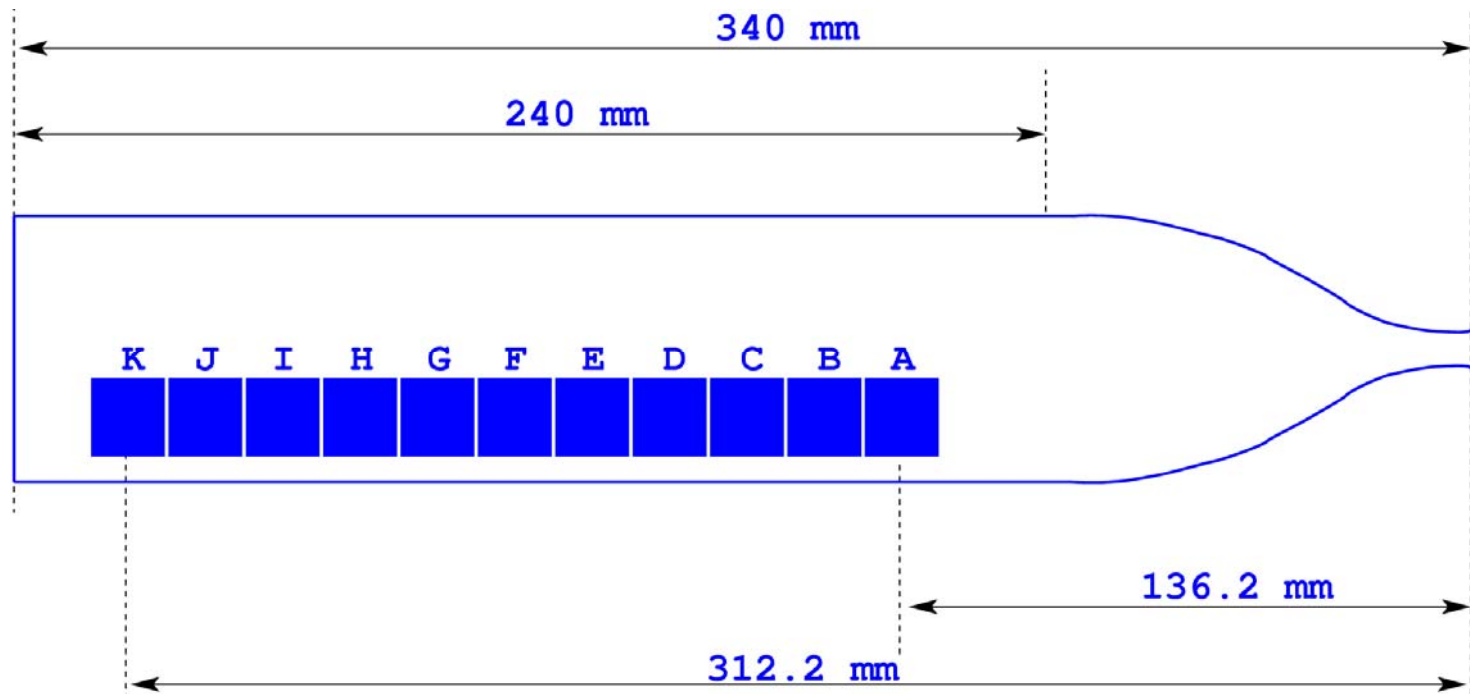
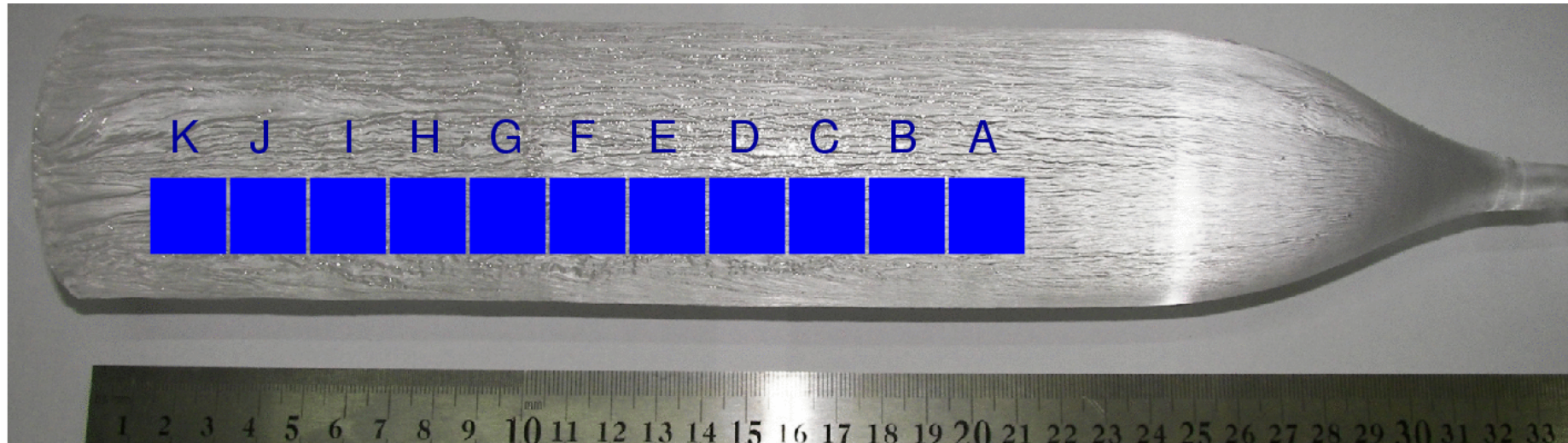
R&D aiming at producing crystals for HEP experiments



Broken after 1st attempt cutting two $2.5 \times 2.5 \times 20 \text{ cm}$ samples



11 Samples from the 1st Ingot

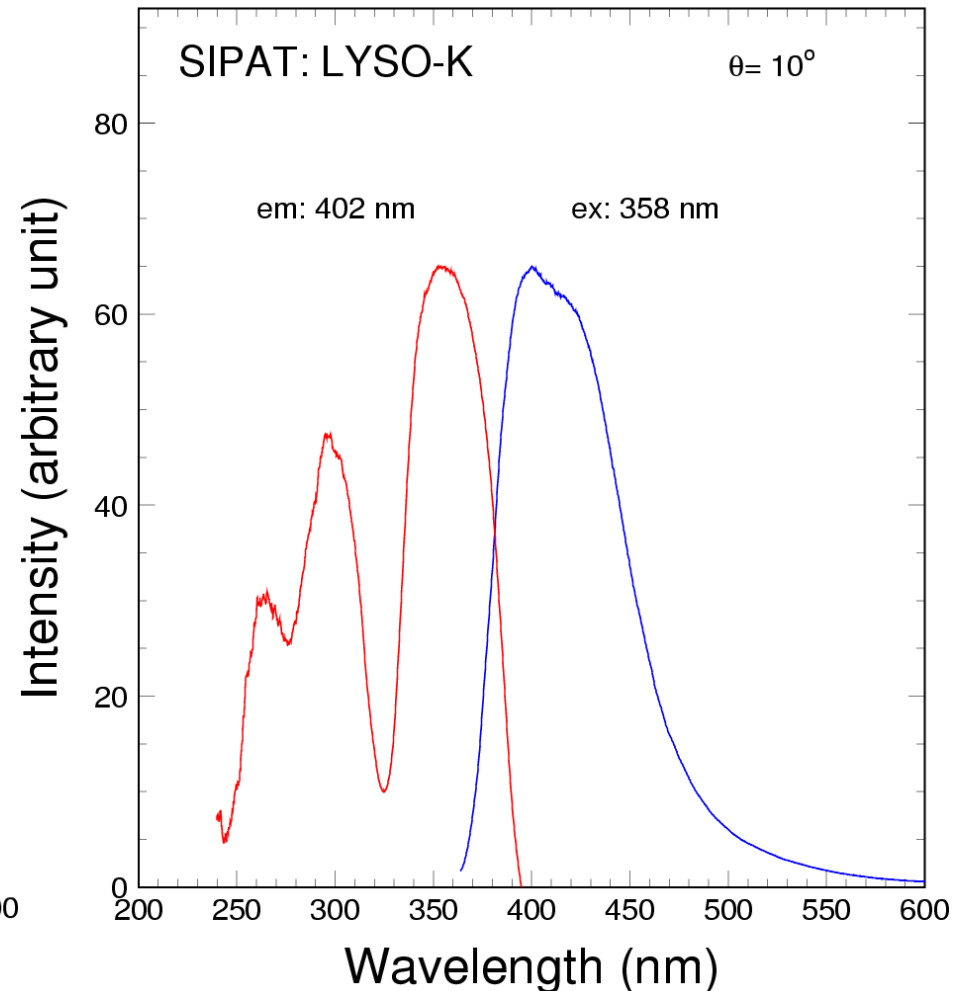
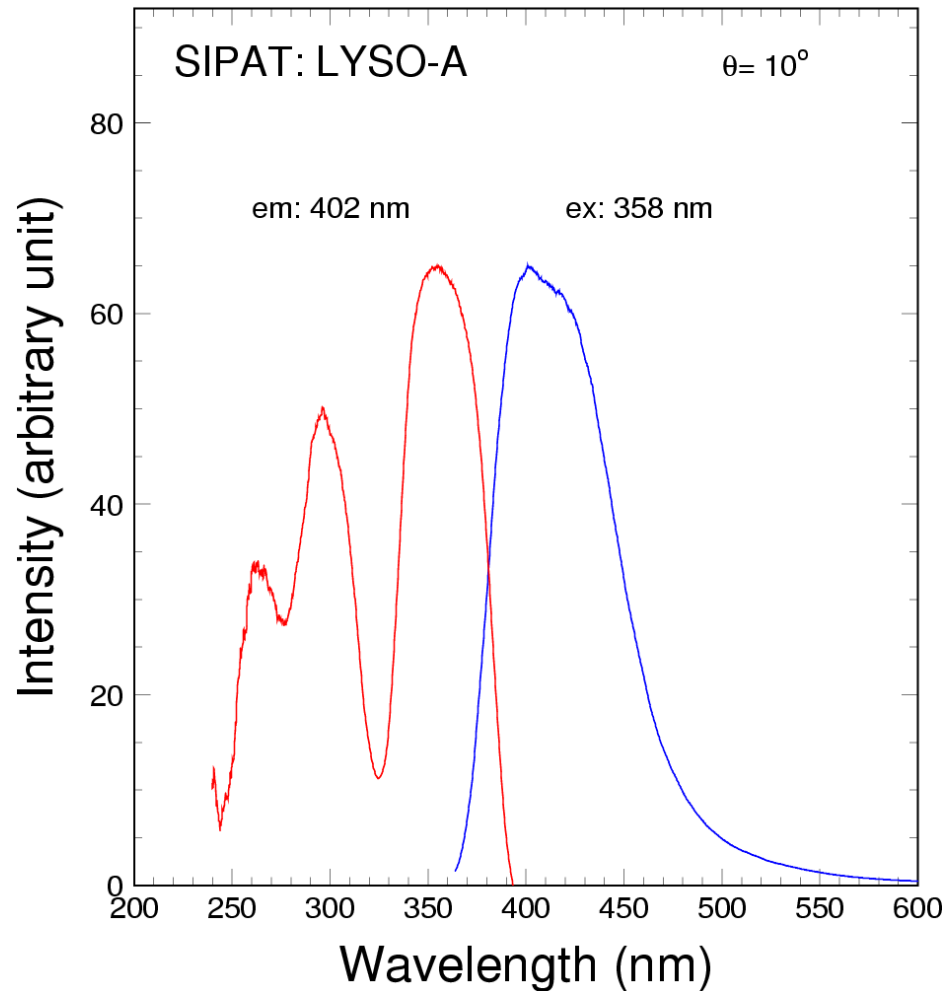




Excitation & Photo-Luminescence



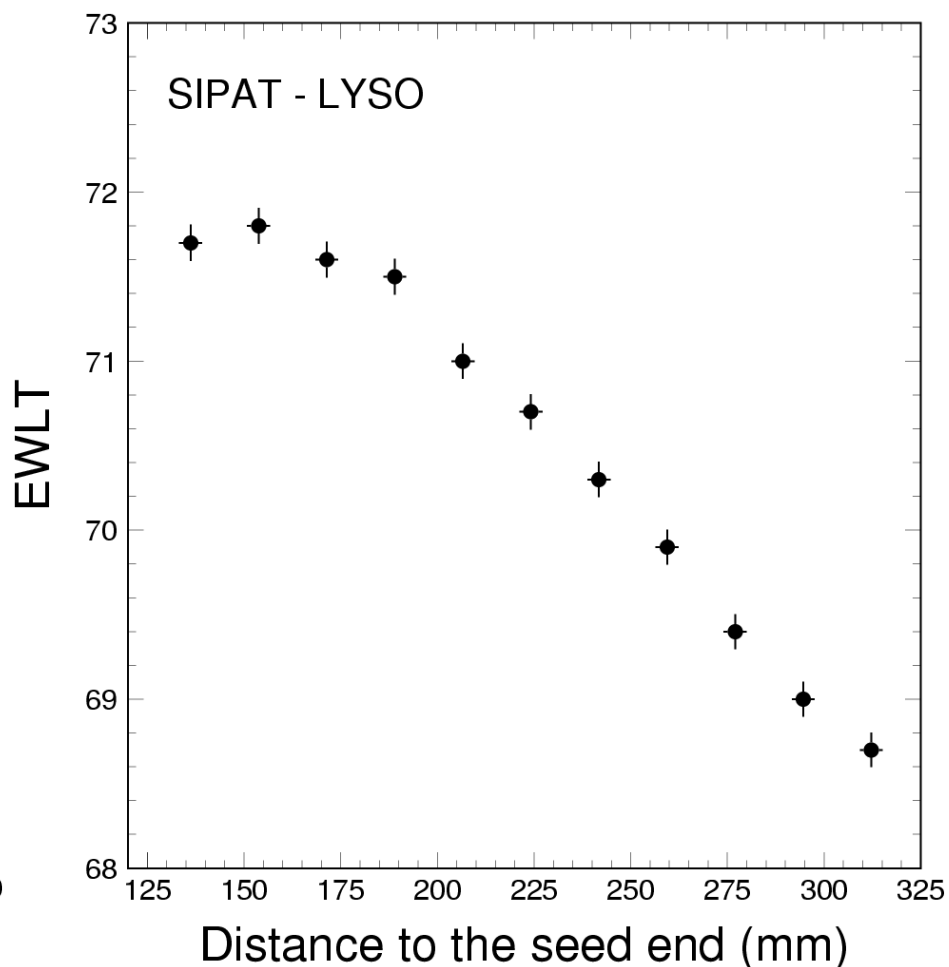
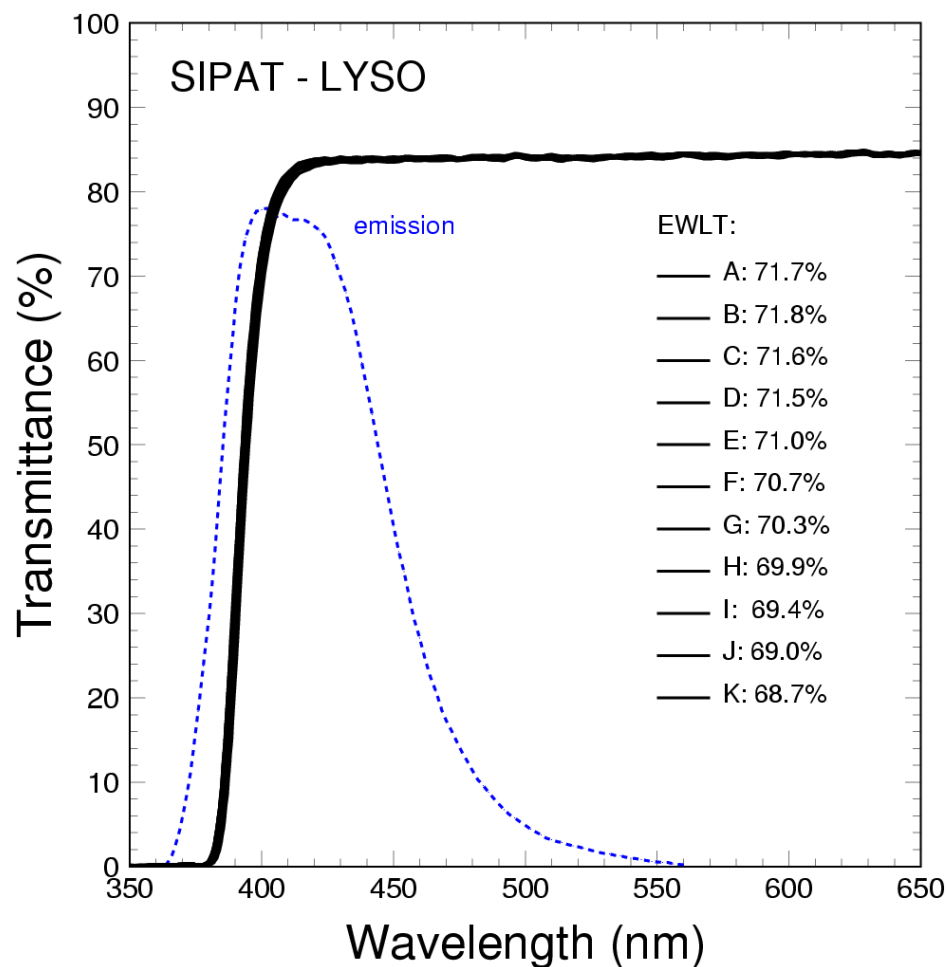
No variation in excitation and emission



Transmittance



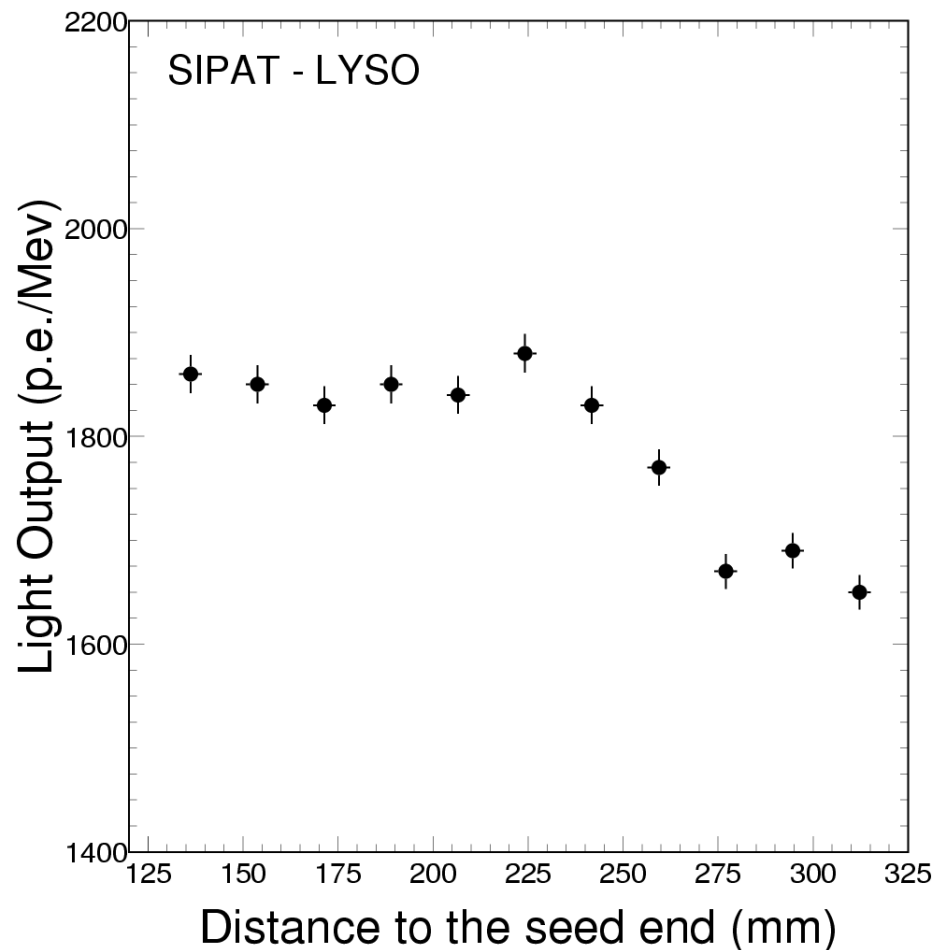
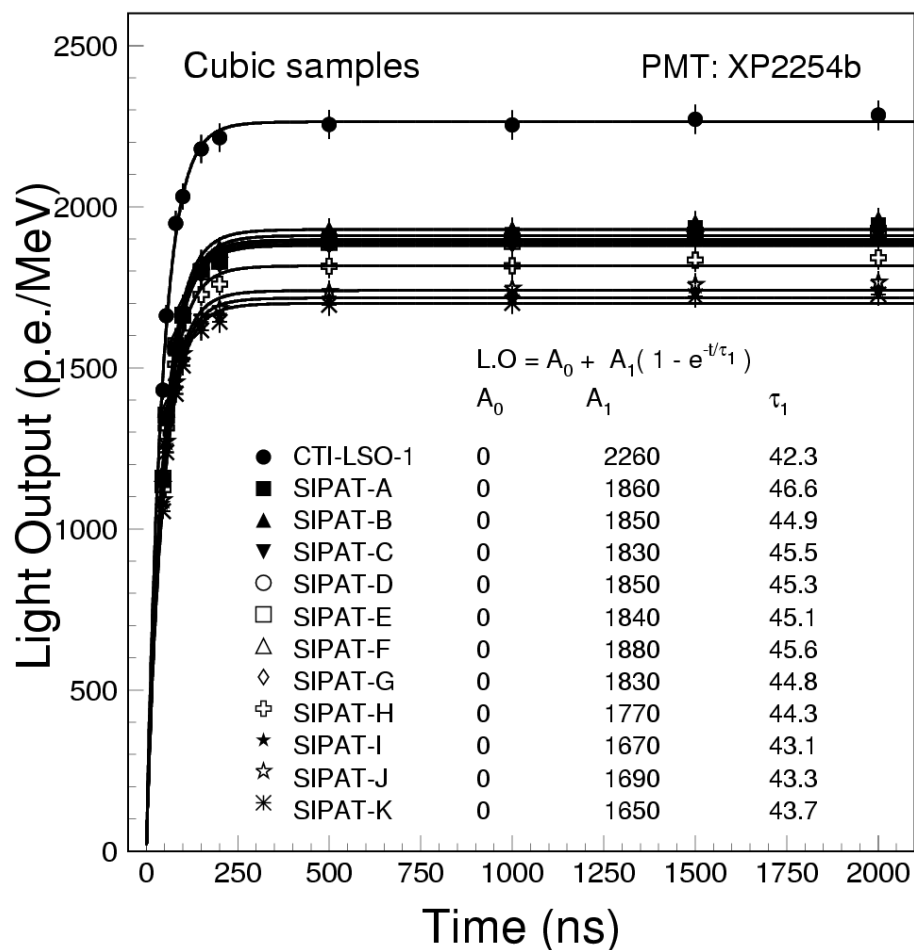
Transmittance degrades after 5 cm from the seed



Light Output



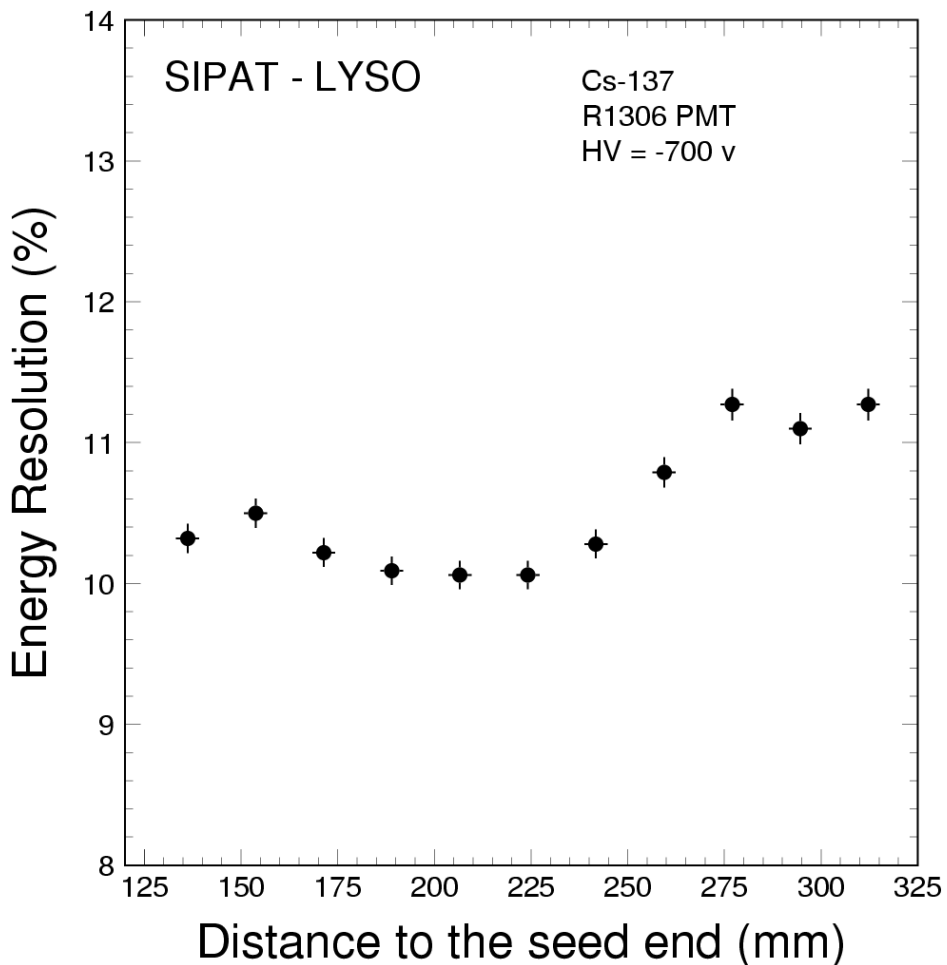
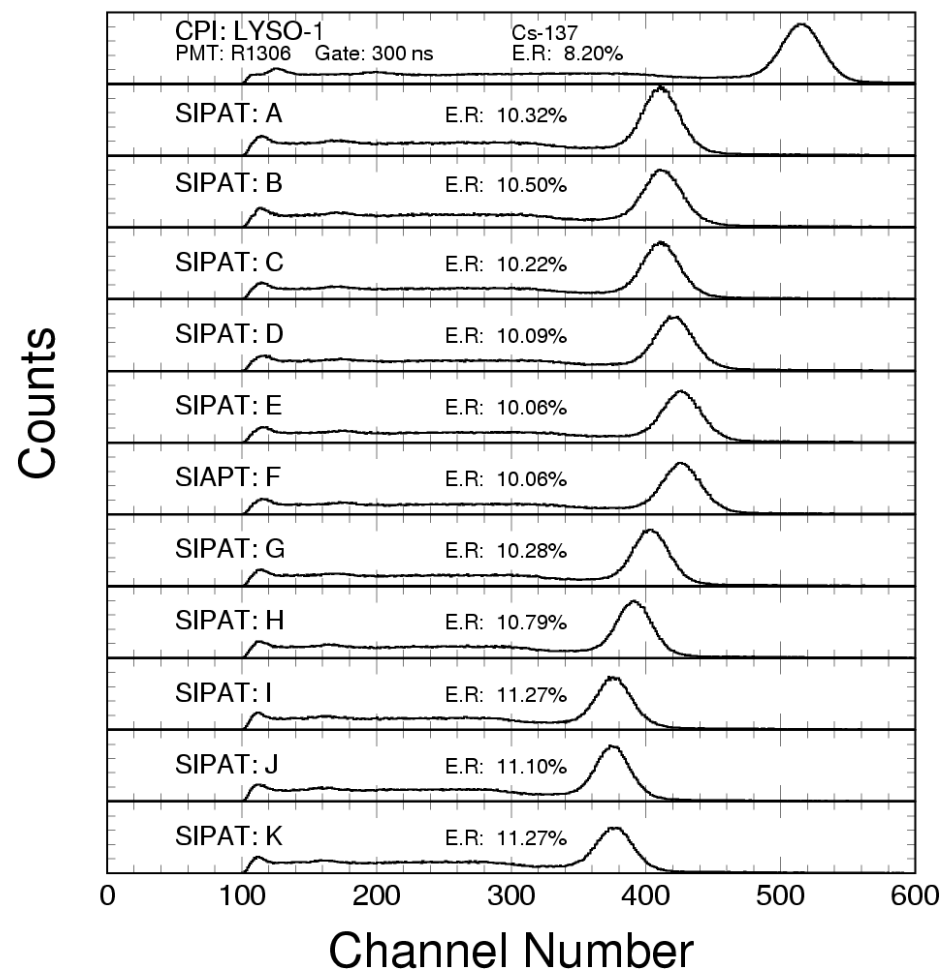
Light output degrades after 10 cm from the seed



Energy Resolution



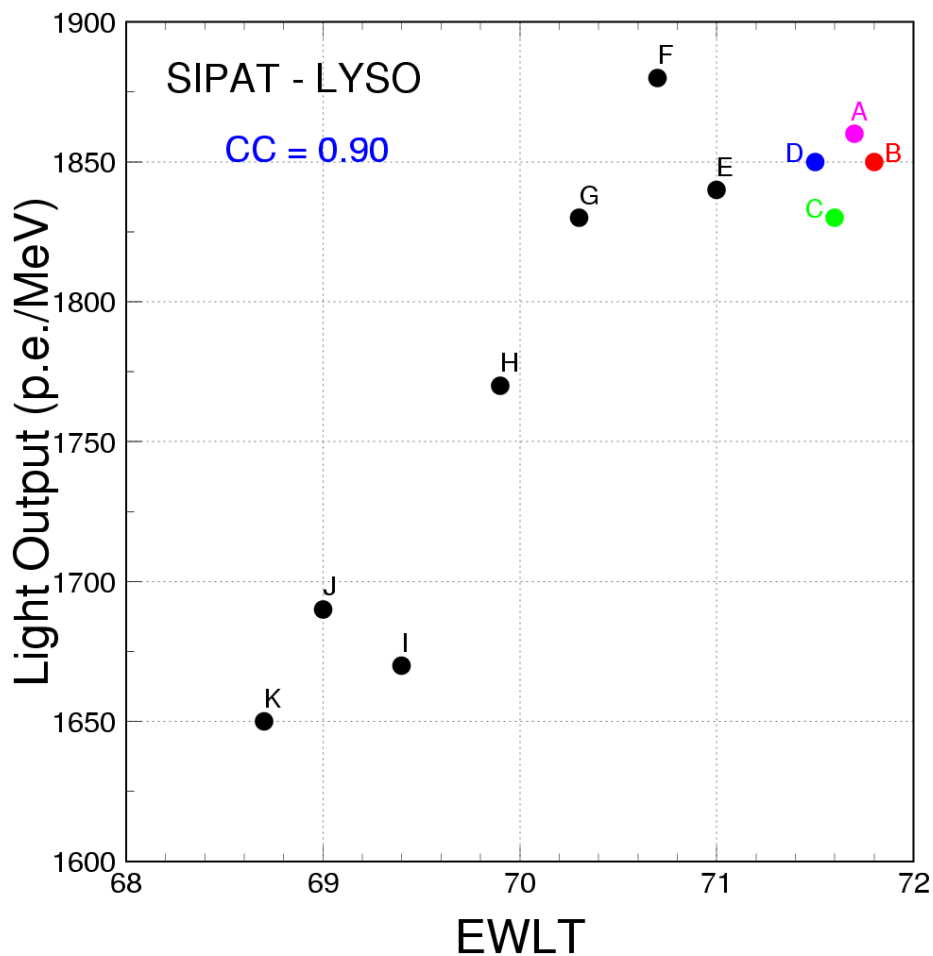
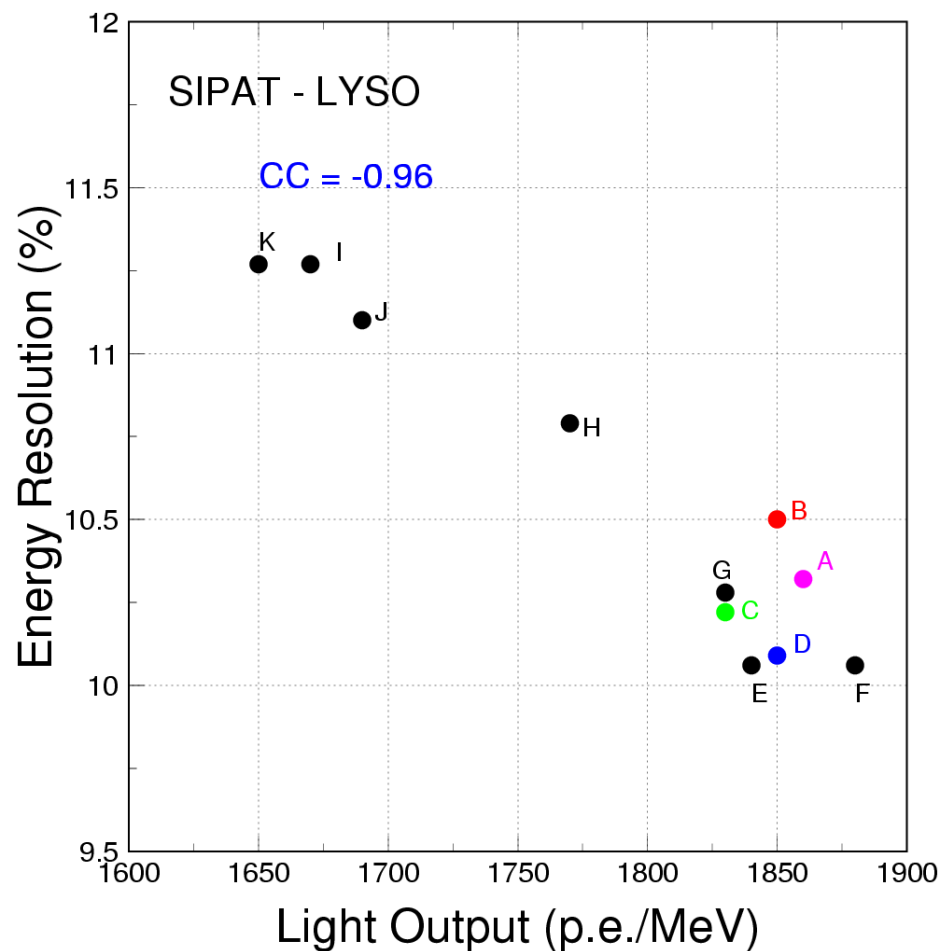
Resolution degrades after 10 cm from the seed



Excellent Correlations



Between light output and resolution/transmittance





Ce & Y Concentrations by GDMS

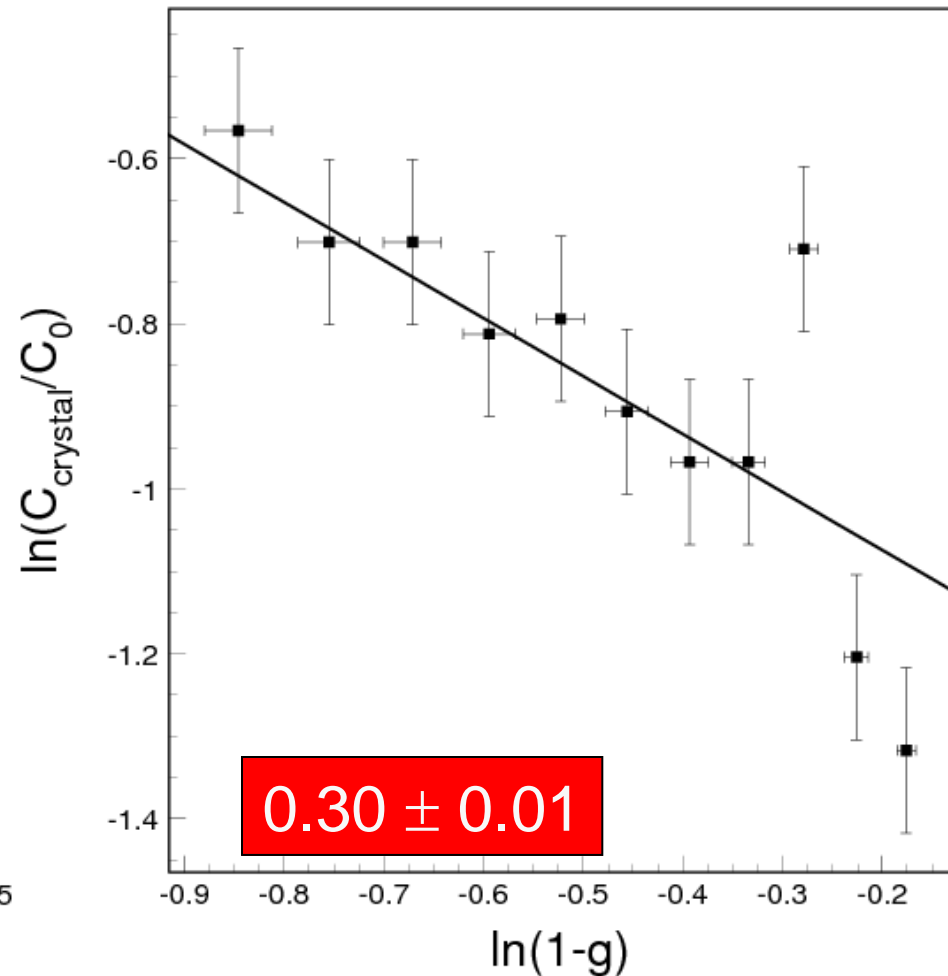
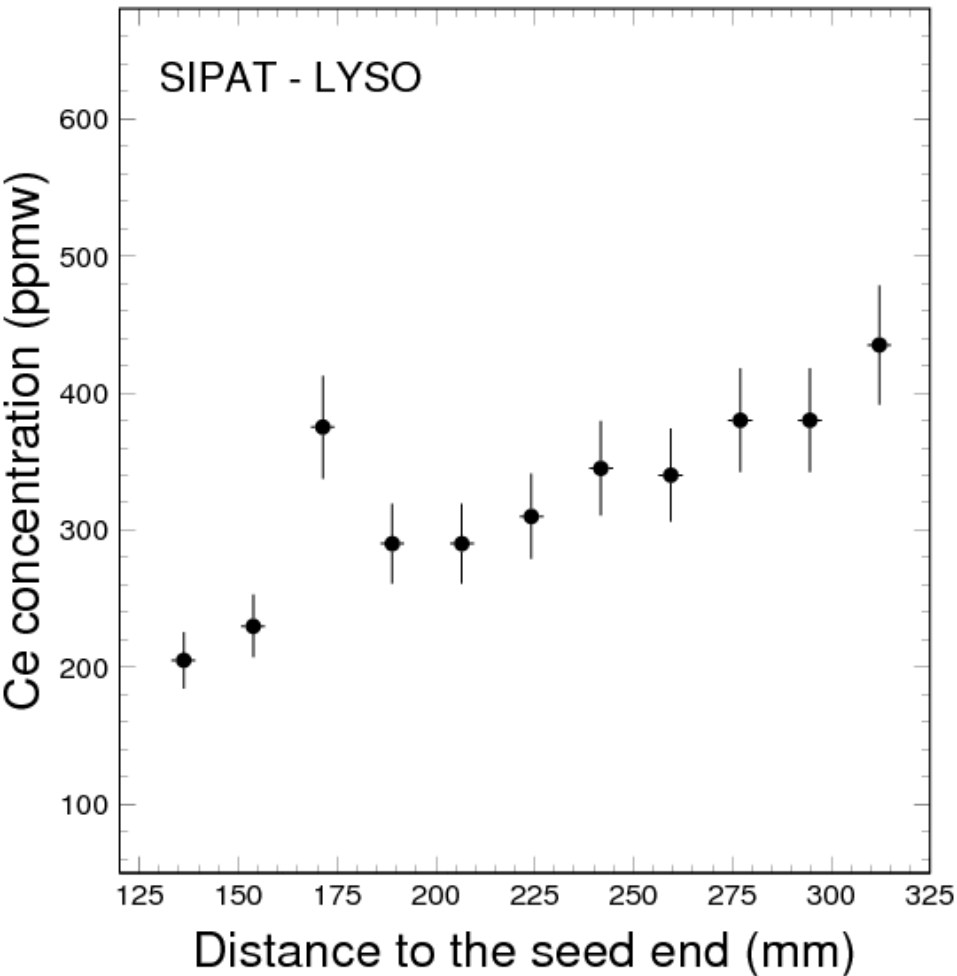


ID	Ce (ppmw)	Y (W%)
SIAPT-A	205	1.11
SIAPT-B	230	1.05
SIAPT-C	375	1.42
SIAPT-D	290	1.32
SIAPT-E	290	1.09
SIAPT-F	310	1.33
SIAPT-G	345	1.43
SIAPT-H	340	1.28
SIAPT-I	380	1.33
SIAPT-J	380	1.24
SIAPT-K	435	1.45

Cerium Segregation in LSO



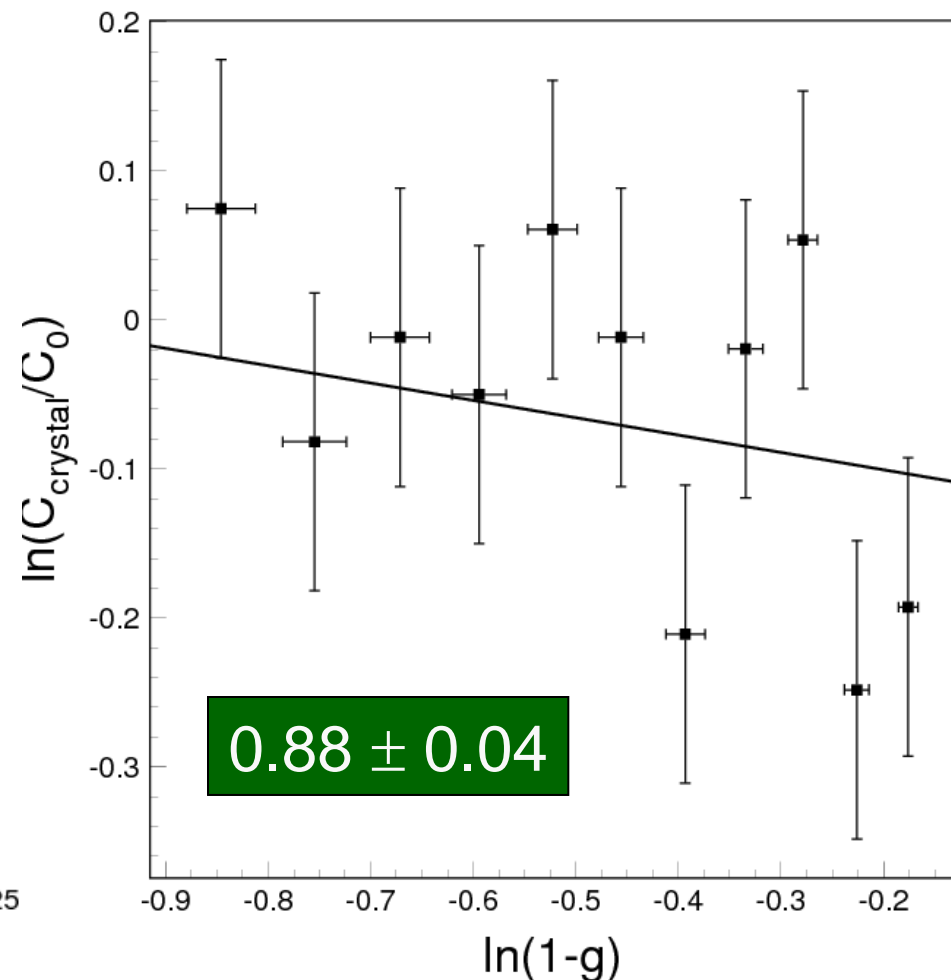
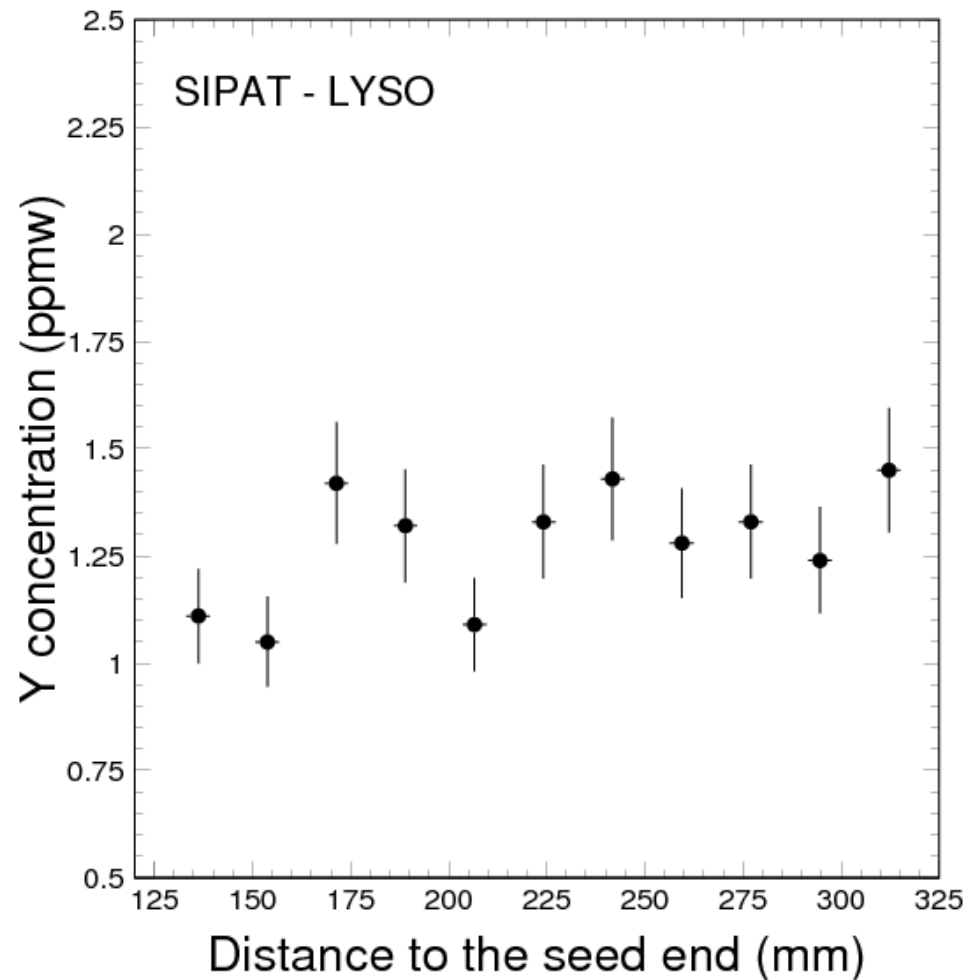
Ce is not easy to enter LSO lattice.
It may be important to make it uniform.



Yttrium Segregation in LSO



Yttrium is more uniform in LSO.
Its distribution seems not a concern.

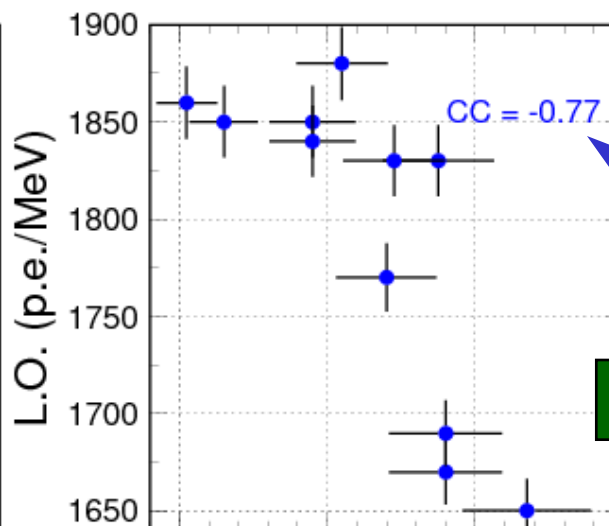
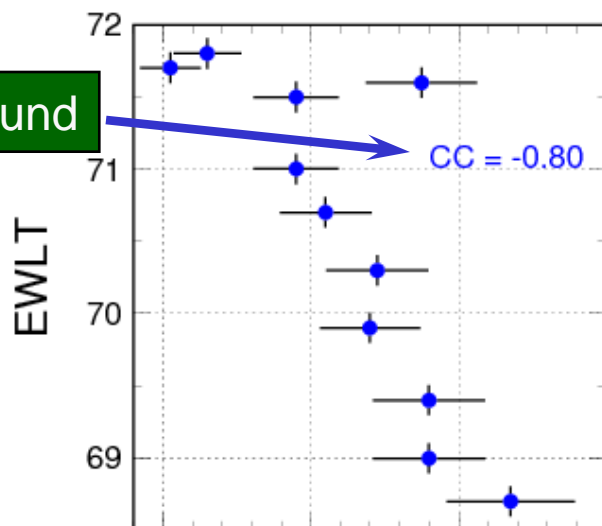




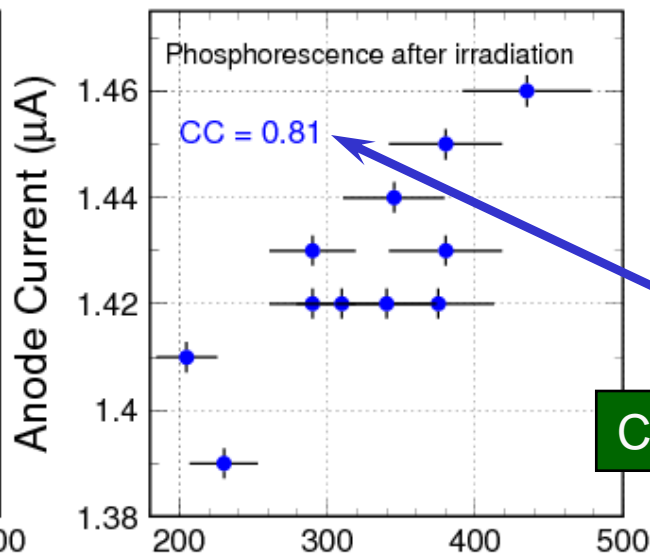
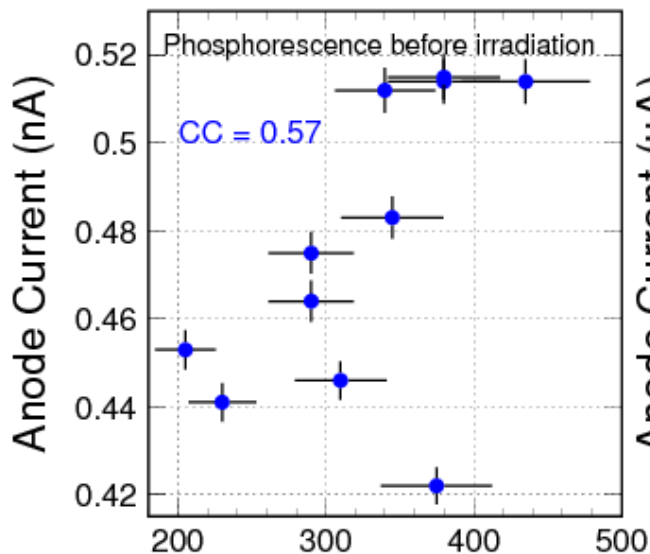
Correlation: Ce and EWLT/L.O./Phors.



Correlation found



Correlation found



Correlation found

Ce Concentration (ppmw)



Summary



- LSO/LYSO crystals are a good candidate for future precision crystal calorimeters.
- Progress has been made in understanding LSO and LYSO crystals.
- Development of cost effective LYSO crystals for SuperB experiments is being pursued.
- Thanks to the DOE ADR program for supporting this work.