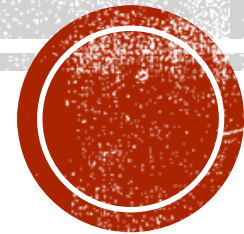


# **AN OVERVIEW OF SCINTILLATORS AND THEIR APPLICATIONS IN HIGH ENERGY PHYSICS**

By Blake D. Leverington



# OUTLINE

- Introduction
- Inorganic Scintillators
  - Excitons and Activators
  - Stokes Shift and the Franck-Condon Principle
- Organic Scintillators
  - The Foerster Resonant Energy Transfer (FRET) Mechanism
- The LHCb Scintillating Fibre Tracker

*“Scintillation is a luminescence induced by ionizing radiation in transparent dielectric media.”*

*Taken from:*

**Inorganic Scintillators for Detector Systems**

Physical Principles and Crystal Engineering

Second Edition

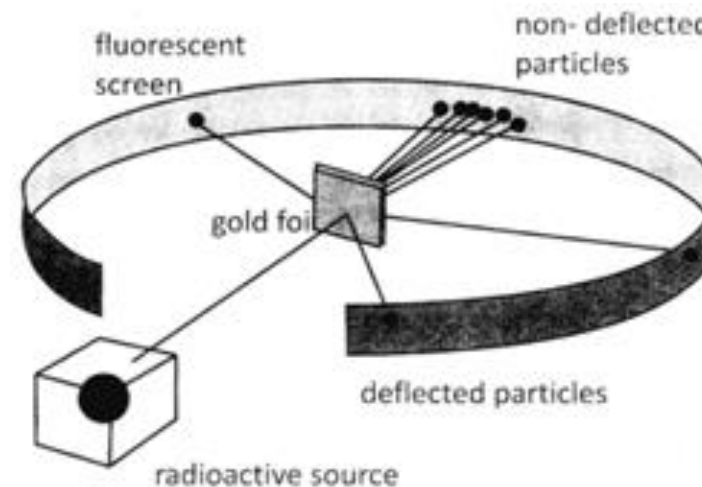
Paul Lecoq • Alexander Gektin • Mikhail Korzhik

- Detectors for radioactive particles

- *The modern era of particle physics began with Rutherford using Zinc Sulfide to observe alpha particles in 1899.*

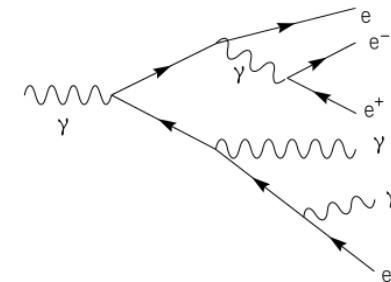
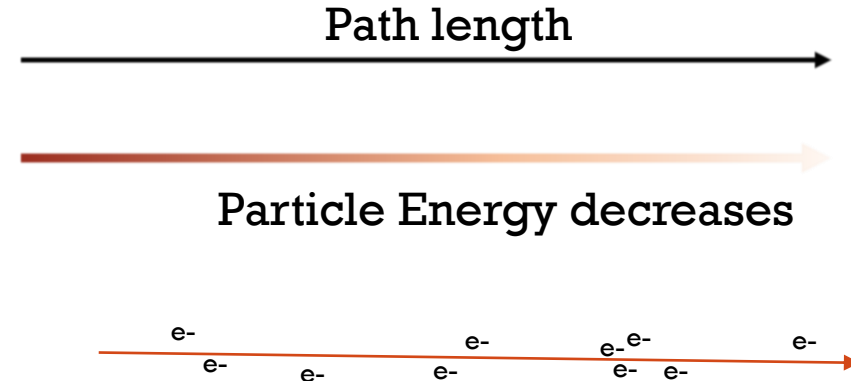


**Ernest Rutherford**  
(1871-1937)



# IONIZATION PROCESS IN MATERIALS

- Neutrons and hadrons
  - Nuclear interactions (pi0 showers, beta or gammas, knock-on electrons)
  - Ionization of atoms (stripped electrons)
- Electrons and photons
  - Bremsstrahlung, pair production (high E) → electromagnetic shower
  - photo-absorption (low to med E)
  - Compton scattering dominates below 1 MeV
- Lots of “hot” electrons produced
  - Convert this to a signal!



# TYPES OF SCINTILLATOR

## ▪ Inorganic Scintillators

- Crystals (hundreds, such as  $\text{PbWO}_4$ )
- High Z atoms
- High density ( $3\text{-}8\text{ g/cm}^3$ )
  - Good for compact calorimeters
- High light yield ( $10\text{-}100\text{ kph/MeV}$ )
  - Good resolution for calorimetry
- Slower ns – us decay times
  - Can be bad for timing resolution
  - afterglow
- Expensive (production)
- Hygroscopic (will reduce light yield)
- Temp dependent Light yield (%/degree)

- Gases (nitrogen + noble gases)
- Glasses (boron silicates)

## ▪ Organic Scintillators

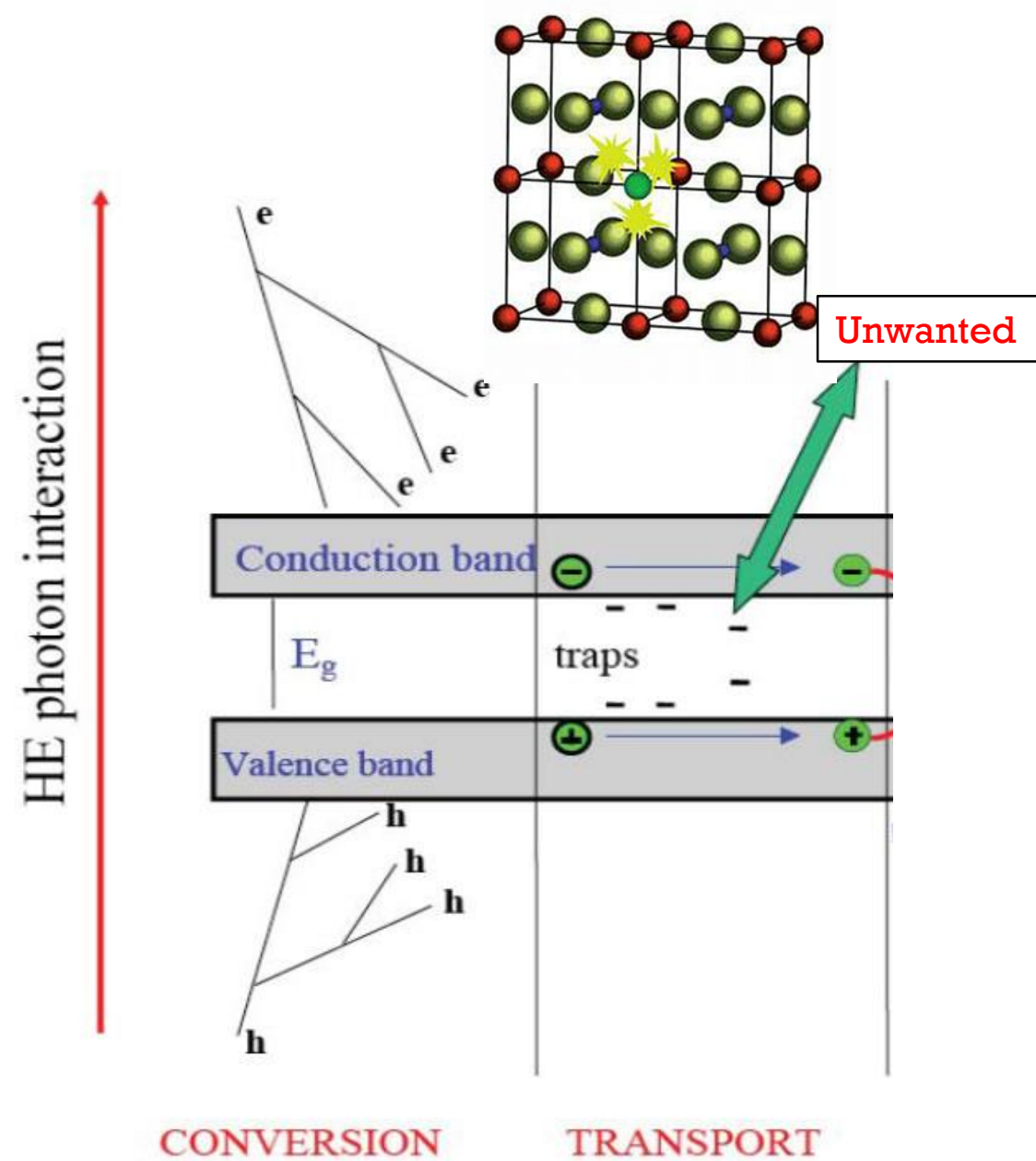
- Crystals (anthracen, naphthalene, stilbene, etc)
- Liquids (in organic solvents, benzene ring needed)
- Plastic (organic fluors in a base polymer)
  
- Lower Z / density ( $1\text{-}2\text{ g/cm}^3$ )
- Lower light yield ( $1\text{-}10\text{ kph/MeV}$ )
- Fast ns decay times
  - Good timing resolution
  
- Cheaper
- Shapeable/machinable
- Independent of temperature ( $-60\text{ -- }+20$ )

# INORGANIC CRYSTALS

- In the case of ordered materials like crystals,  $< \text{keV}$  electrons begin to couple with the lattice

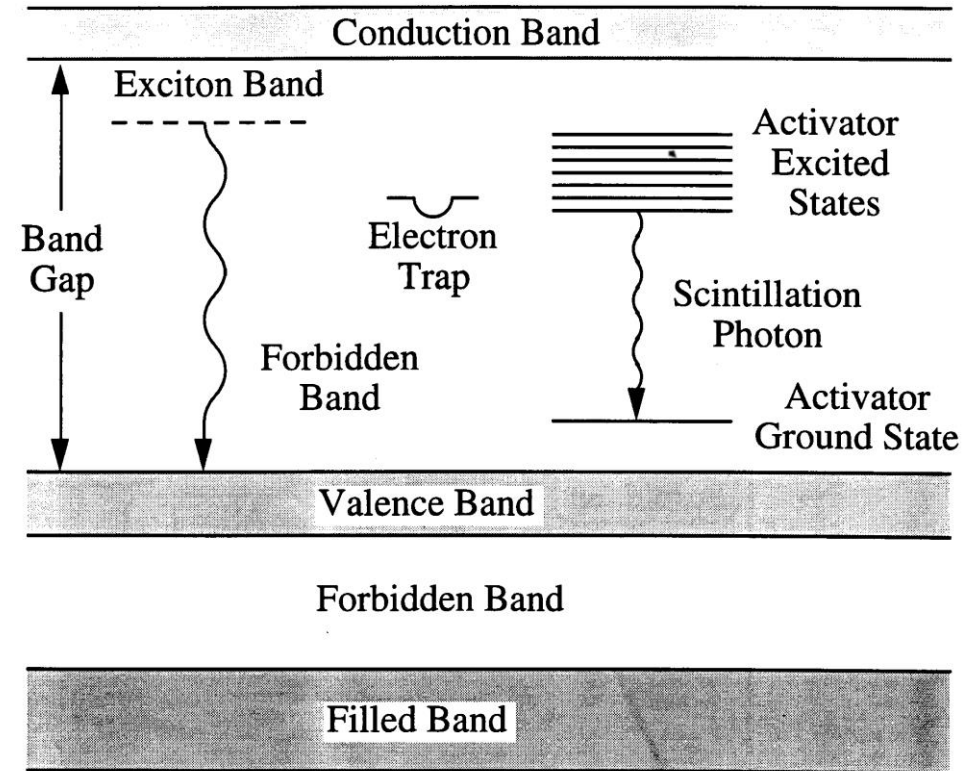
## Mobility of electrons

- Some electrons are ionized and free ( $E > E_{\text{ion}}$ )
- Some electrons are excited from the valence band to the **conductance band** (or just above and relax) ( $E > \sim E_g$ )
- Some loosely bind with a hole and **form an exciton** ( $E < \sim E_g$ )



# INORGANIC SCINTILLATOR

- The crystal must contain luminescent centers to be a scintillator
  - Intrinsic
  - Extrinsic



- Beware of traps (defects) – reduces conversion efficiency, delays emission
- Once the electrons or energy can travel, high probability of absorption by a luminescent center
  - **Relaxes to its ground state -> photon emission**



# INORGANIC SCINTILLATOR

- The crystal must contain luminescent centers to be a scintillator

- Intrinsic

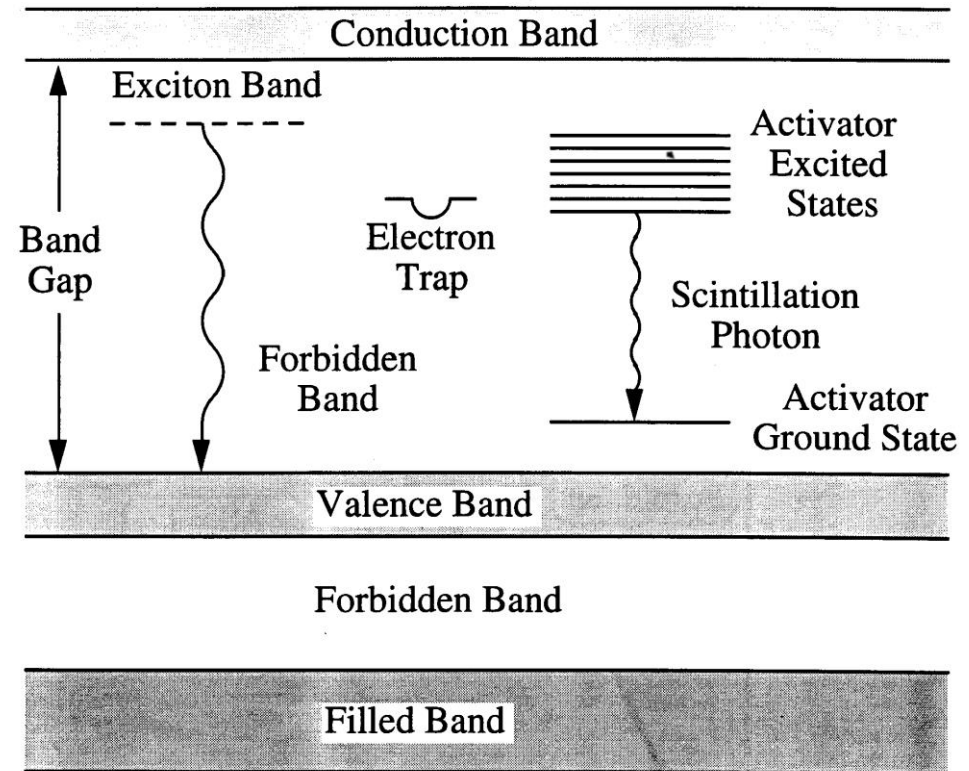
- “involve electron-hole recombination; free, **self-trapped (PbW04)**, and defect-trapped exciton luminescence; constituent transition group or post-transition group ion fluorescence; core-valence band transitions; or charge transfer transitions within a molecular complex” –Derenzo NIM 2002
- Internal defects allow for exciton recombination (eg. BGO)

- Extrinsic

- Defects and Impurities
- **Doped with ions (activator)**
- Combination of activator ions and with impurities

- Beware of traps (defects) – reduces conversion efficiency, delays emission

- Once the electrons or energy can travel, high probability of absorption by a luminescent center
  - **Relaxes to its ground state -> photon emission**

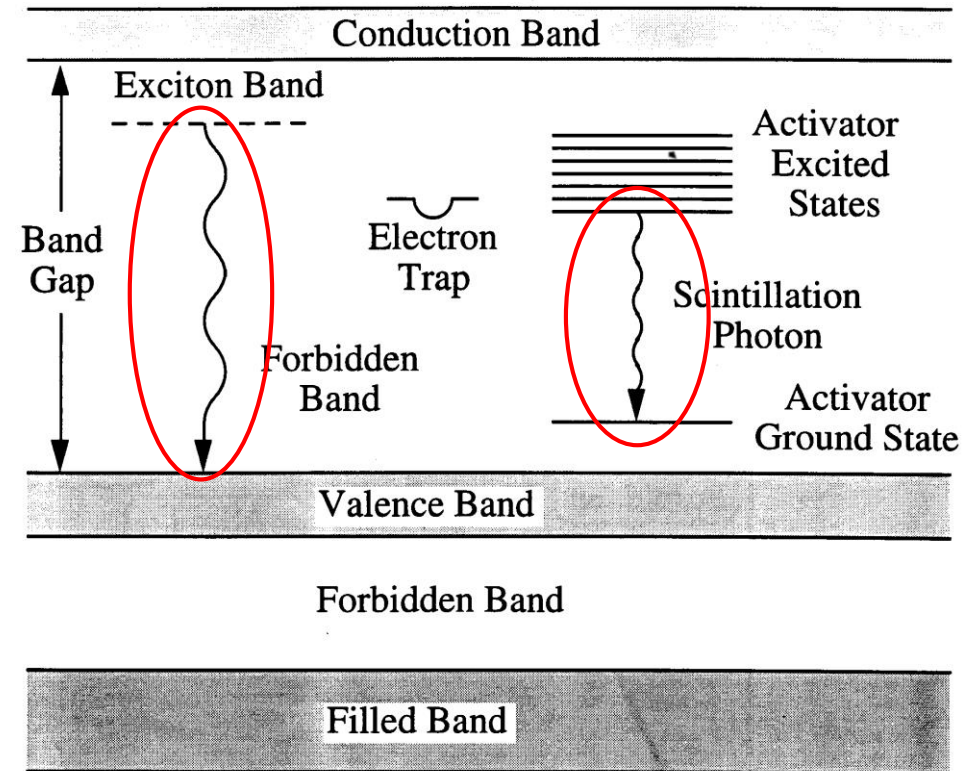


S.E. Derenzo, *Scintillation Counters, Photodetectors and Radiation Spectroscopy*, IEEE Short Course Radiation Detection and Measurement, 1997 Nuclear Science Symp.

# INORGANIC SCINTILLATOR

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    - Internal defects allow for exciton recombination (eg. BGO)
  - **Extrinsic**
    - Defects and Impurities
    - Doped with ions (activator)
    - Combination of activator ions and with impurities
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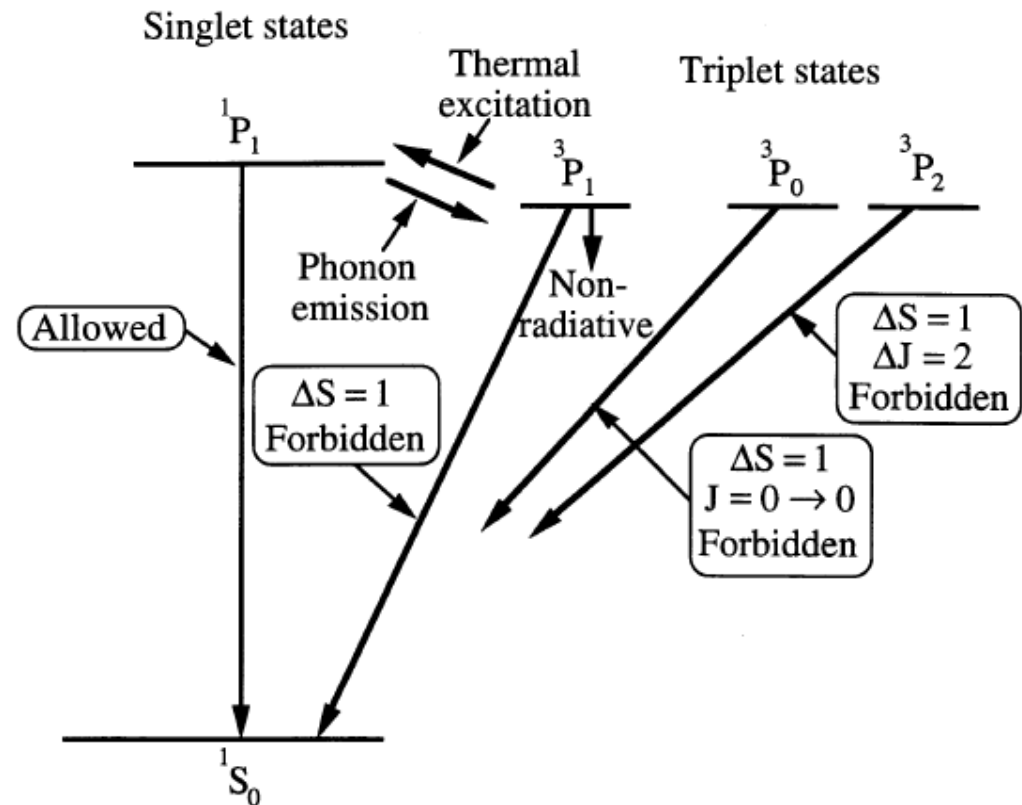
An **exciton** is a loosely bound (0.01 eV) e-h pair that can move within the crystal; integer spin



S.E. Derenzo, *Scintillation Counters, Photodetectors and Radiation Spectroscopy*, IEEE Short Course Radiation Detection and Measurement, 1997 Nuclear Science Symp.

# LUMINESCENCE

- Time to excited state is very fast
  - (femtoseconds)
- Radiative emission from Singlet states
  - Fluorescence
  - Fast (nanoseconds)
- Radiative emission from Triplet states
  - Phosphorescence
  - Slow (microseconds)
- Quenching will compete with these processes
  - Relaxation to ground state through vibrational modes



# EXAMPLE SCINTILLATOR

$$LY \left( \frac{ph}{MeV} \right) = N_{e-h} S Q = \left( \frac{10^6}{\beta E_g} \right) S Q$$

$N_{e-h}$  = number of  $e - h$  pairs

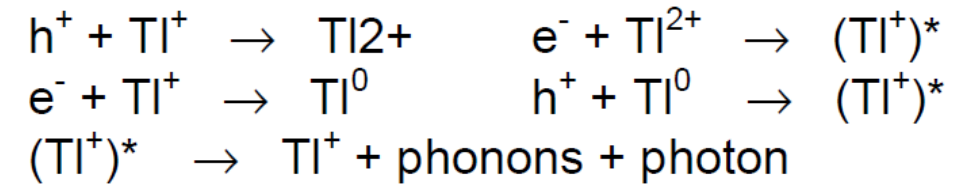
$E_g$  = band gap energy

$\beta$  = ionisation energy conversion eff.  
(typically  $\sim 2-7$ )

$S$  = carrier transfer efficiency  
(least well known: material, temperature, impurity dependent)

$Q$  = *q. e. of luminescent center*

NaI(Tl):



*Introduction to Radiation Detectors and Electronics*  
III. Scintillation Detectors

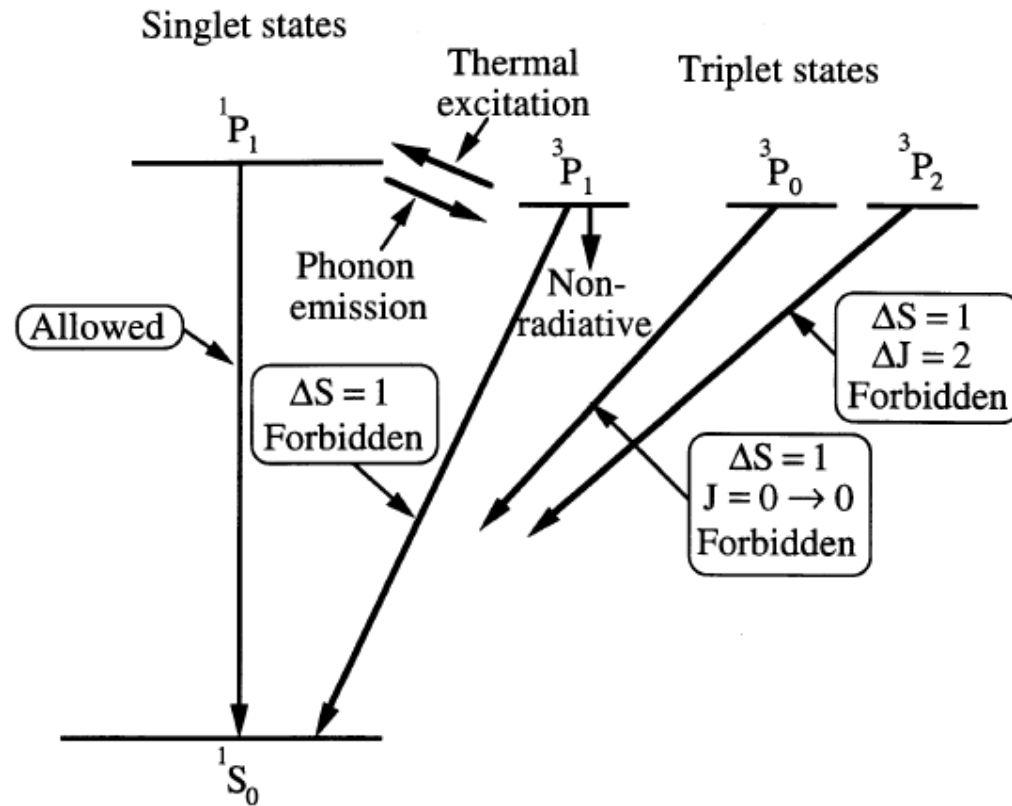
Copyright © 1998 by Helmuth Spieler

(NaI(Tl)) has a light output  $\sim 40,000$  photons per MeV of energy deposit.

- $Q(Tl^+) \sim 1$
- Rise time is fairly fast ( $\tau_R \sim 10$  ns).
  - Slow rise times are from slow carriers
- the decay time is rather slow ( $\tau_D \sim 250$  ns).
  - some transitions are forbidden (phosphorescence)

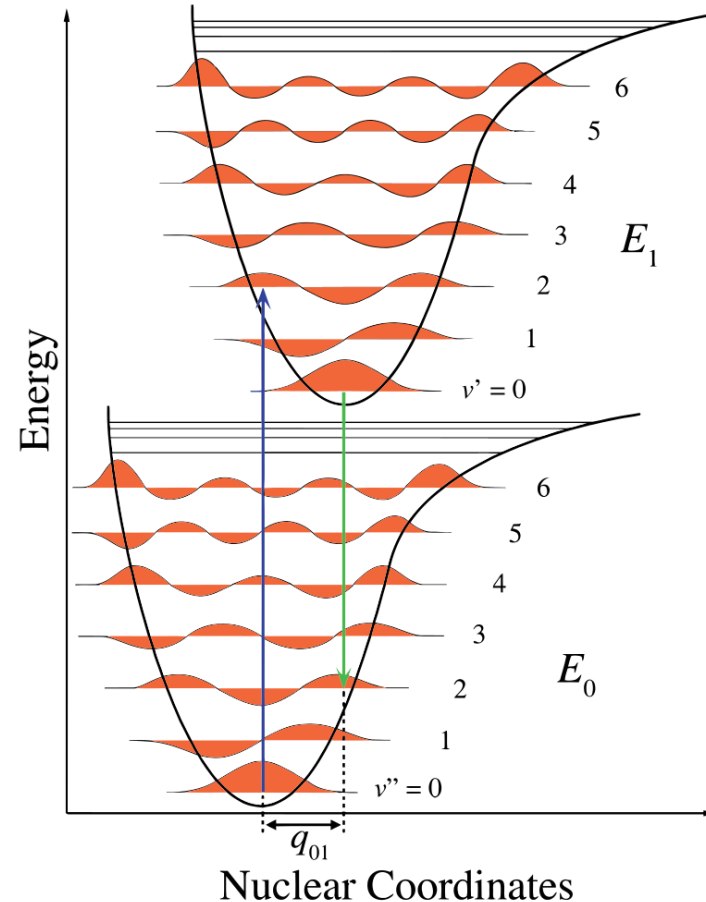
# WHY ARE THE PHOTONS NOT REABSORBED INSTANTLY?

Photon re-absorption?



# STOKES SHIFT AND THE FRANCK-CONDON PRINCIPLE

- Franck-Condon principle (semi-classical)
  - The electronic transitions are much faster than the nuclear motions
  - new vibrational level must be instantaneously compatible with the nuclear positions and momenta of the vibrational level of the molecule
  - transitions between vibrational sublevels  $\nu=0$  and  $\nu=2$  and  $\nu=1 \rightarrow \nu=0$  are favoured over  $\nu=0 \rightarrow \nu=0$
- $h\nu < E_{\text{abs}}$  (Stokes Shift) = longer wavelength

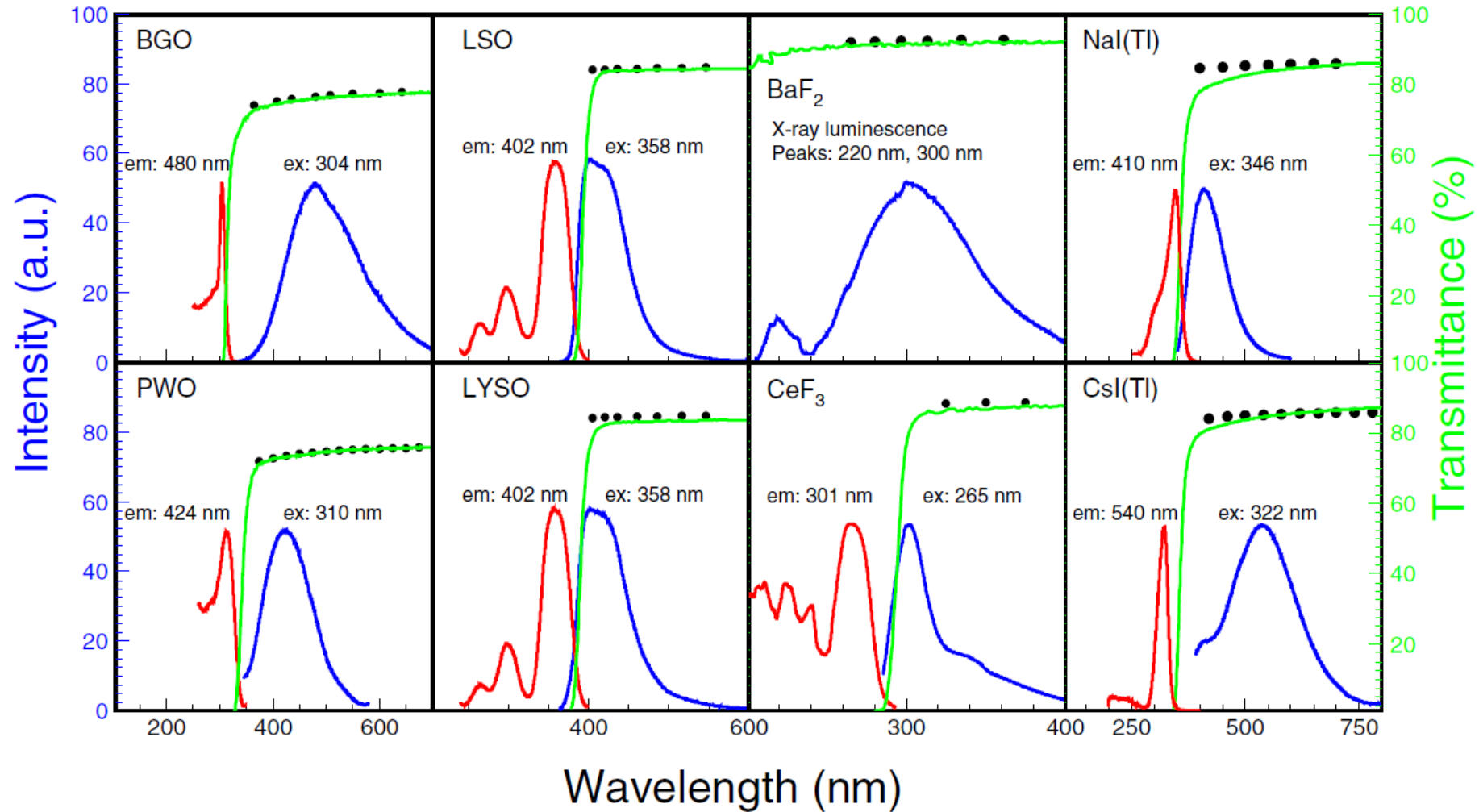


Excitation  $E_{0,0} \xRightarrow{fs} E_{1,2}$

Relaxation  $E_{1,2} \xRightarrow{ps} E_{1,0}$

Radiative  $E_{1,0} \xRightarrow{ns} E_{0,2}$

# Inorganic Scintillator Absorption and Emission spectra & Transmission



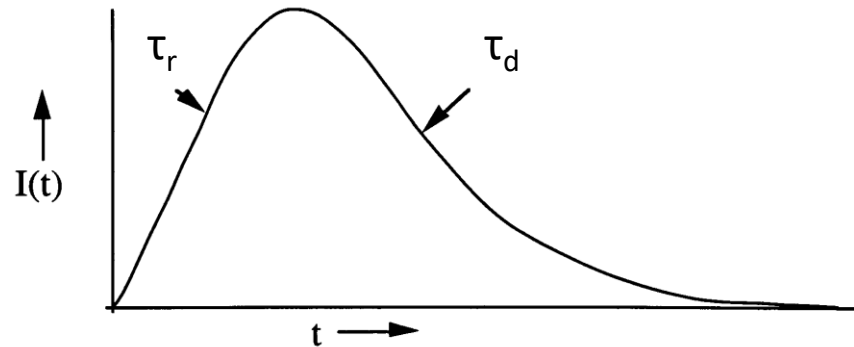
2007 IEEE Nuclear Science Symposium Conference Record N49-1, Rihua Mao et al.

# TIMING (SIMPLE APPROXIMATION)

- Rise time,  $\tau_r$ , due to transfer time from vibration modes to luminescent centres
- Decay time,  $\tau_d$ , due to emission and quenching of luminescent centres

- Timing resolution dependent on the arrival time,  $t_{1st}$ , of the first photon

$$I(t) = I_0(e^{-t/\tau_d} - e^{-t/\tau_r})$$



$$t_{1st} = \tau_d / N_{ph} \text{ for } \tau_r < \tau_d / N_{ph}$$

$$t_{1st} = \sqrt{2\tau_r\tau_d / N_{ph}} \text{ for } \tau_r > \tau_d / N_{ph}$$

From "Inorganic Scintillators for Detector Systems," Paul Lecoq

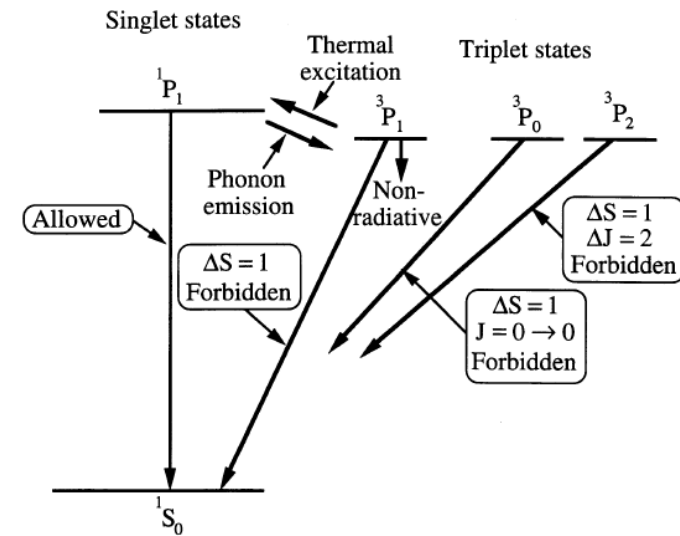
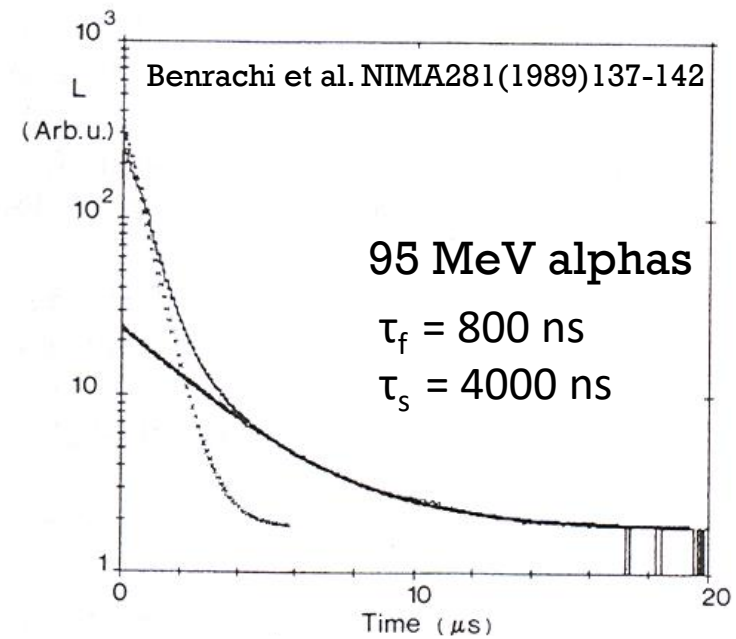


# PARTICLE ID

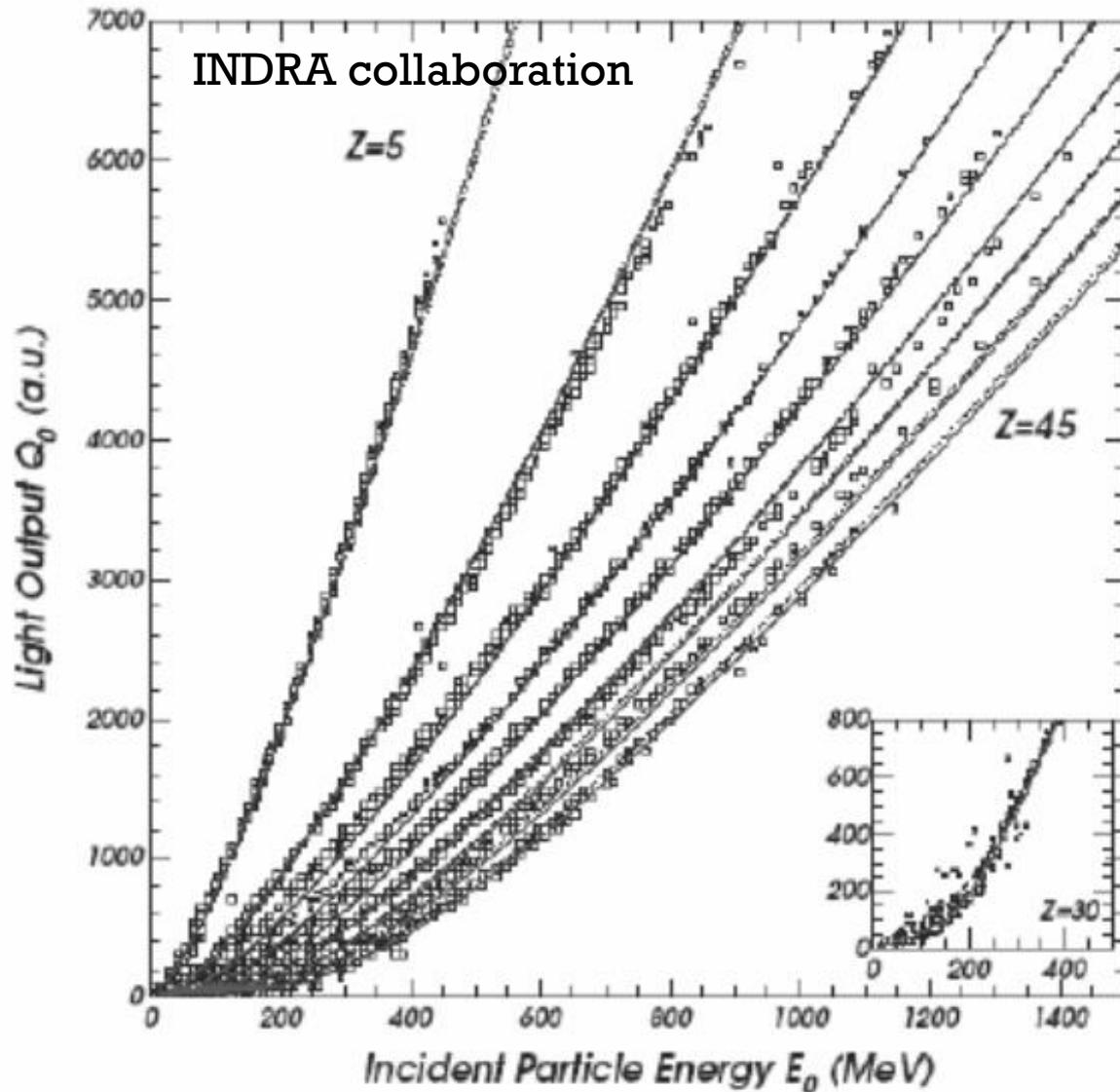
- For CsI(Tl), the pulse shape has two distinct components, fast and slow

$$I(t) = \frac{h_f}{\tau_f} e^{-\frac{t}{\tau_f}} + \frac{h_s}{\tau_s} e^{-\frac{t}{\tau_s}}$$

- $\tau_f$  is dependent on ionization density
  - Excitation to states with forbidden decays more likely
- $\tau_s$  is independent
- $R = \frac{h_s}{(h_s+h_f)}$  increases with ionization density



# Inorganic Scintillator CsI(Tl)



Example: n- $\gamma$  discrimination

P. Sperr, H. Spieler, M.R. Maier, NIM 116(1974)55

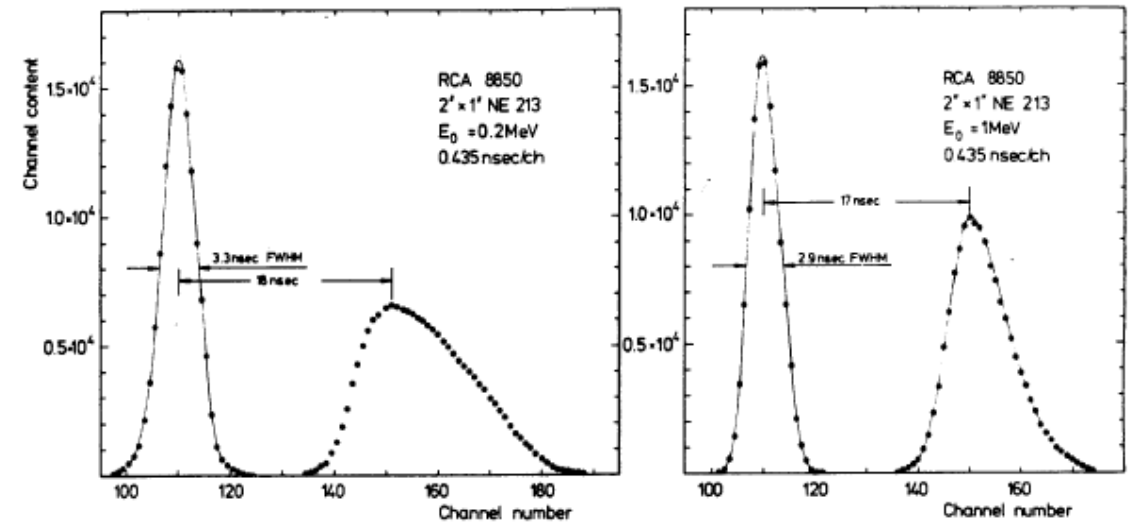


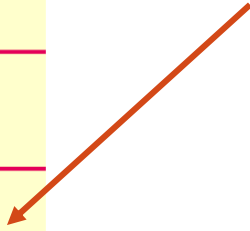
Fig. 5. Neutron-gamma timing distributions with a small scintillator (2" diam. x 1") for two threshold energies.

# OTHER RELEVANT PROPERTIES

- Refractive index:
  - Coupling efficiency to photodetectors
  - High (1.5-2)
- Transmission of emitted photons
  - Optical transparency of the crystal
- Density:
  - Compactness = cost \* volume
- Small Moliere radius (high density, lower Z)
  - $RM \sim X_0 (Z + 1.2)/37.74$ .
  - Smaller EM shower radius = better separation
- Radiation and Interaction length
  - Shower containment improves energy resolution
- Temperature dependence
- Wavelength matching to photodetector quantum efficiency

Crystal	NaI(Tl)	CsI(Tl)	CsI	BaF <sub>2</sub>	BGO	LYSO(Ce)	PWO
Density (g/cm <sup>3</sup> )	3.67	4.51	4.51	4.89	7.13	7.40	8.3
Melting Point (°C)	651	621	621	1280	1050	2050	1123
Radiation Length (cm)	2.59	1.86	1.86	2.03	1.12	1.14	0.89
Molière Radius (cm)	4.13	3.57	3.57	3.10	2.23	2.07	2.00
Interaction Length (cm)	42.9	39.3	39.3	30.7	22.8	20.9	20.7
Refractive Index <sup>a</sup>	1.85	1.79	1.95	1.50	2.15	1.82	2.20
Hygroscopicity	Yes	Slight	Slight	No	No	No	No
Luminescence <sup>b</sup> (nm) (at peak)	410	550	310	300 220	480	402	425 420
Decay Time <sup>b</sup> (ns)	245	1220	26	650 0.9	300	40	30 10
Light Yield <sup>b,c</sup> (%)	100	165	3.7	36 4.1	21	85	0.3 0.1
d(LY)/dT <sup>b</sup> (%/°C)	-0.2	0.4	-1.4	-1.9 0.1	-0.9	-0.2	-2.5

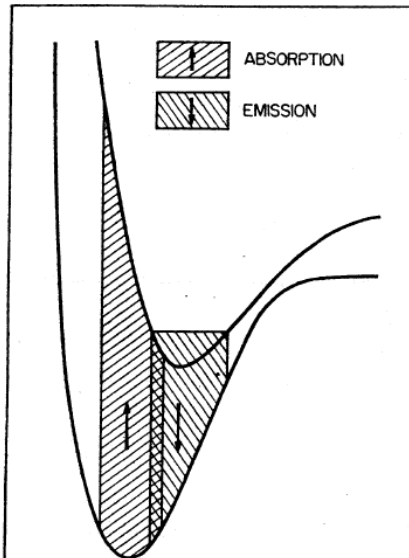
Complex interaction of luminescent centres results in multiple decay time components



A more extensive HTML table is available at: <http://scintillator.lbl.gov>

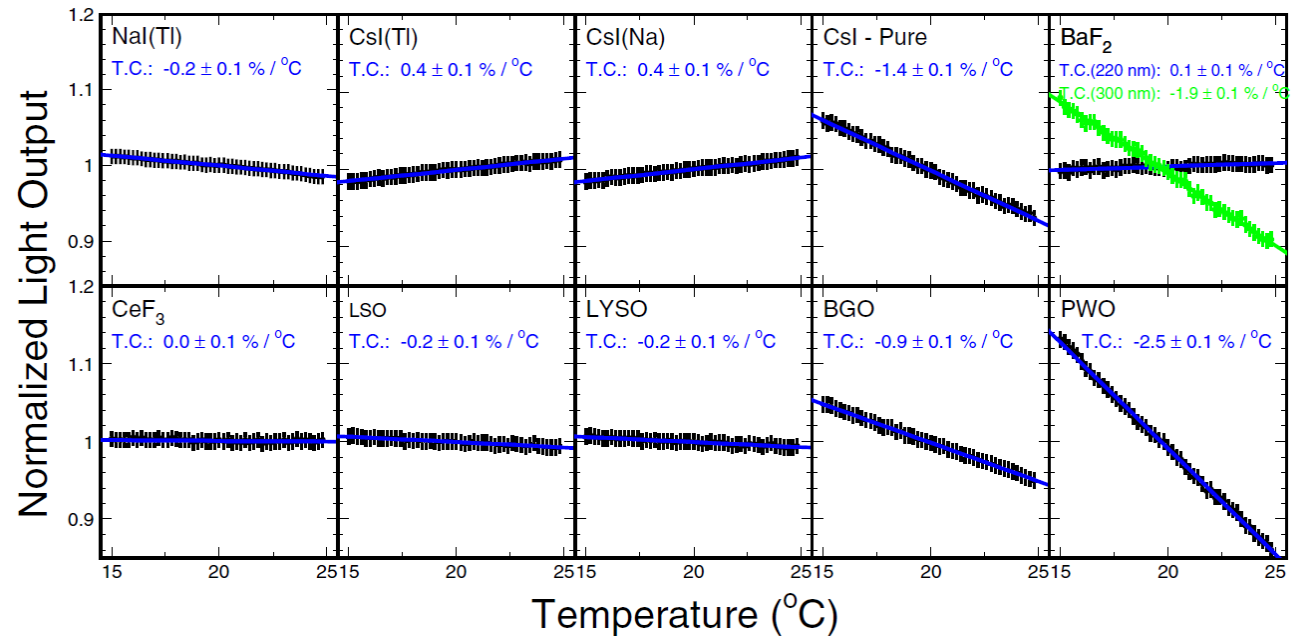
# TEMPERATURE DEPENDENCE

- At higher temperatures, the vibrational states widen
- Simple case: Overlapping ground states result in reabsorption (quenching) and loss of light yield



- Improves the timing, however

J.B. Birks, *The Theory and Practice of Scintillation Counting*, New York, 1964



2007 IEEE Nuclear Science Symposium Conference Record N49-1, Rihua Mao et al.

# ENERGY RESOLUTION

- Mass resolution through 2 photon decay channel (i.e. H->gamma gamma)
- Want  $\partial E/E = 0.5\%$  for 100 GeV photons
- $a=2\%$  is achievable for modern homogenous EM calorimeters

$$\frac{\delta E}{E} \sim \frac{1}{\sqrt{N}}$$

- $B < 0.5\%$  is hard
  - Need  $X_0 \sim 25$  for 100 GeV photons
  - The shower is contained in the readout cluster
  - Temperature dependence can increase  $b_F$ ; also production variation

$$\frac{\delta_M}{M} = \frac{1}{2} \left[ \frac{\delta E_{\gamma_1}}{E_{\gamma_1}} \oplus \frac{\delta E_{\gamma_2}}{E_{\gamma_2}} \oplus \frac{\delta\theta}{\tan(\theta/2)} \right]$$

granularity  $\downarrow$

$$\delta E_\gamma / E_\gamma = a / \sqrt{E} \oplus b \oplus c / E$$

statistical  $\downarrow$   $a / \sqrt{E}$      $\uparrow$   $b$     noise  $\downarrow$   $c / E$   
 constant  $\uparrow$

$$b^2 = b_L^2 \oplus b_F^2 \oplus b_c^2$$

Shower leakage  $\swarrow$   $b_L^2$     Non-uniformity  $\swarrow$   $b_F^2$     Inter-calibration  $\swarrow$   $b_c^2$

From "Inorganic Scintillators for Detector Systems," Paul Lecoq

# DETECTORS WITH INORGANIC SCINTILLATOR

## High Energy Physics

- CMS (LHC):
  - 76150 PbWO<sub>4</sub> crystals (80 tons, 11m<sup>3</sup>)
  - Depth of 25 X<sub>0</sub>
  - 40ns shaping (CMS TDR)
  - 45 mrad/ $\sqrt{E}$  angular precision
  - Resolution of 1-2%
- Babar (SLAC)
  - 6580 CsI(Tl) crystals(6m<sup>3</sup>)
  - Depth of 17 X<sub>0</sub>
  - 270ns integration
- Many other calorimeters...

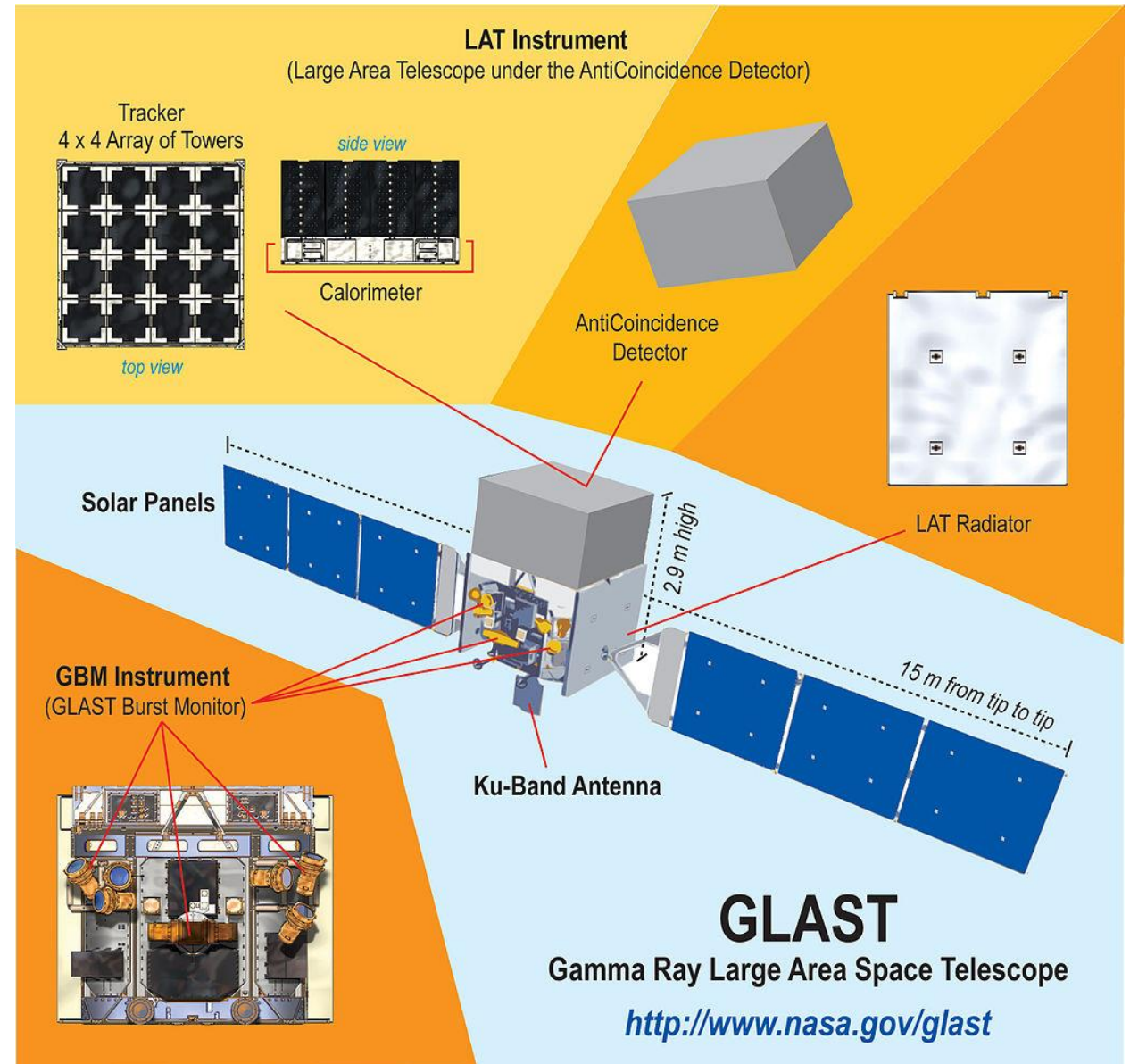


Lead tungstate crystals used in the CMS ECAL.  
<http://cms.web.cern.ch/news/crystal-calorimeter>

# ASTROPHYSICS

## Fermi Gamma-ray Space Telescope (FGST) -- formerly GLAST

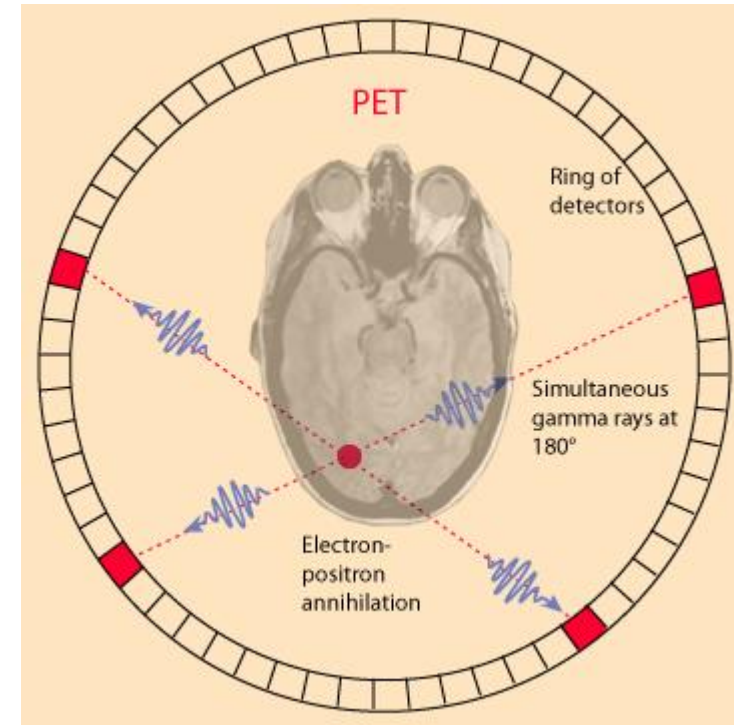
- silicon microstrip detector + calorimeter
- 1536 CsI(Tl) crystals ( $0.3\text{m}^3$ )
- 20 MeV – 300 GeV dynamic range
- 20% energy resolution





# MEDICAL APPLICATIONS

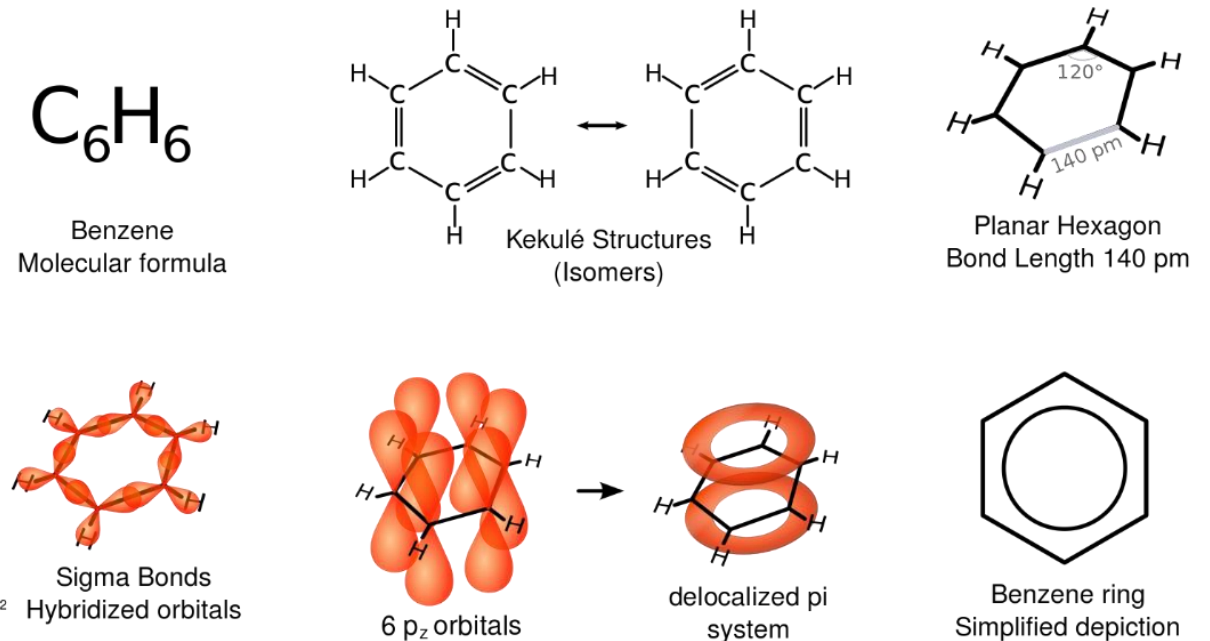
- **PET detectors**
  - BGO, LSO, LYSO crystals
  - Requires  $< \text{ns}$  timing resolution
    - Reduces signal to noise ratio
  - Also good E resolution  $\rightarrow M(e^+ + e^-)$
- X-ray, CT, etc.



From <http://hyperphysics.phy-astr.gsu.edu/hbase/NucEne/nucmed.html>

# ORGANIC SCINTILLATORS

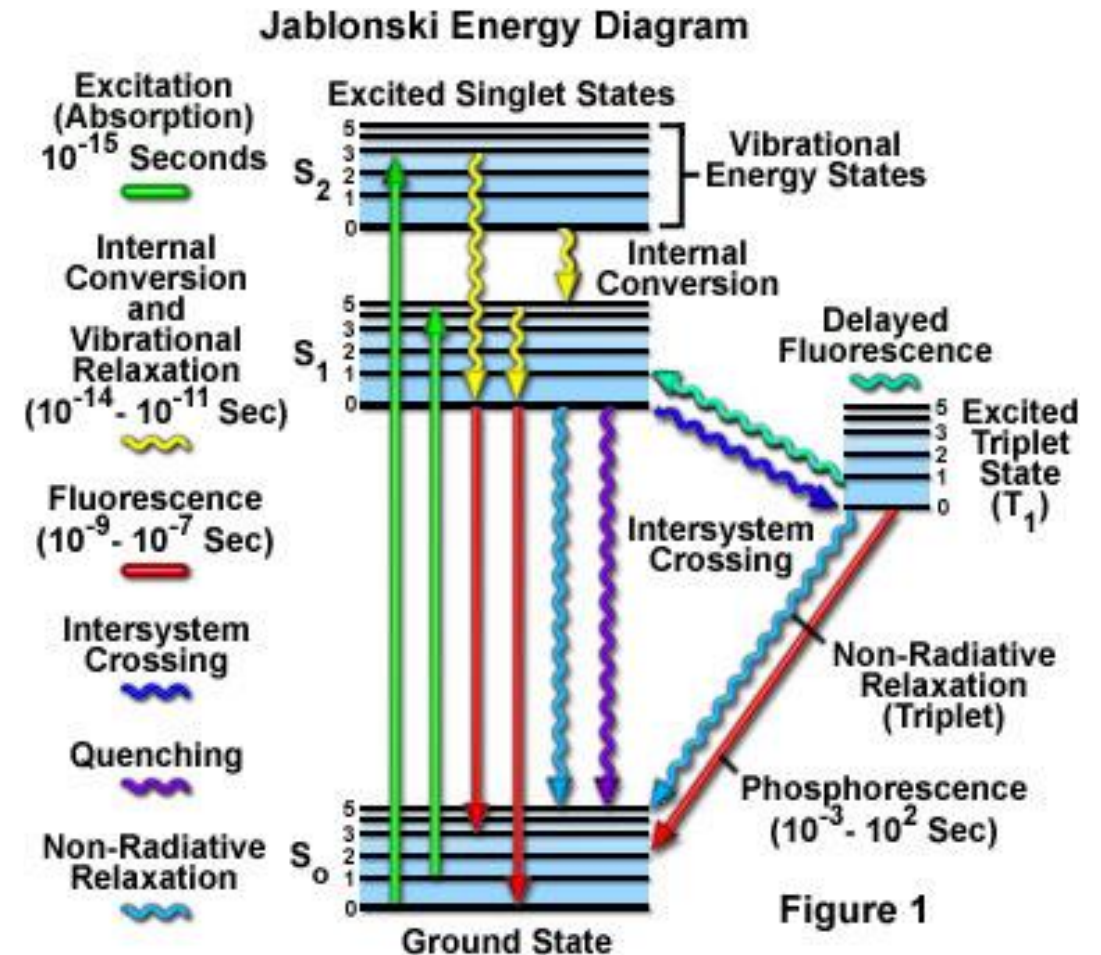
- contain aromatic molecules
  - i.e. they have a **benzene ring**
  - scintillation results from the physics of the benzene ring coupled to the molecule
- $\sigma$ -bonds are in-plane with a bond angle of  $120^\circ$ , from  $sp^2$  hybridization
- $\pi$ -orbitals are out-of-plane; the  $\pi$ -electrons overlap and are completely delocalized
- Scintillation light is produced from the de-excitation of the molecule



By Vladsinger - Own vector drawing based on layout of en:File:Benzol trans.png., CC BY-SA 3.0, <https://commons.wikimedia.org/w/index.php?curid=7536474>

# FLUORESCENCE

- Ionisation energy excites molecule to  $S_{1N}$  or higher ( $\tau = \text{fs}$ )
  - Relaxes to  $S_{10}$  ( $\tau = \text{ps}$ )
- Can decay radiatively to  $S_{0N}$  state ( $\tau = \text{ns}$ )
  - Stokes shift as a result of Franck-Condon principle
  - Relaxes to  $S_{00}$  state
- Overlap with the ground state results in non-radiative decays (quenching)
- $T_1 \leftrightarrow S_0$  forbidden by spin/parity ( $\tau > \text{ms}$ )



<http://micro.magnet.fsu.edu/primer/techniques/fluorescence/fluorescenceintro.html>

# TYPES OF SCINTILLATOR

## Crystals:

- Anthracene : highest light output for organic scint.
  - 17,400 photons/MeV
- Stilbene, TPB, etc.
- good for n/gamma separation
- Expensive!

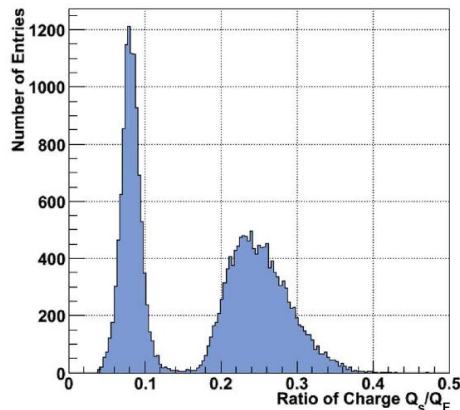
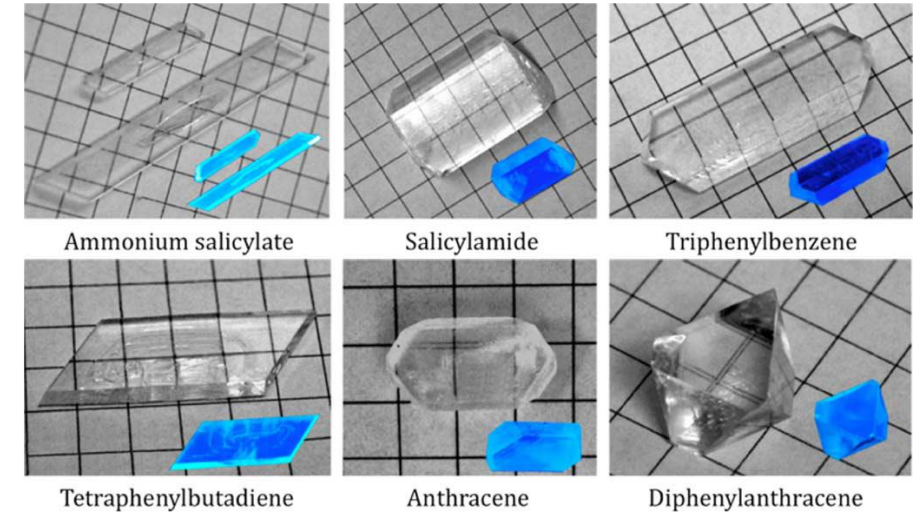
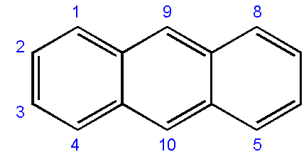
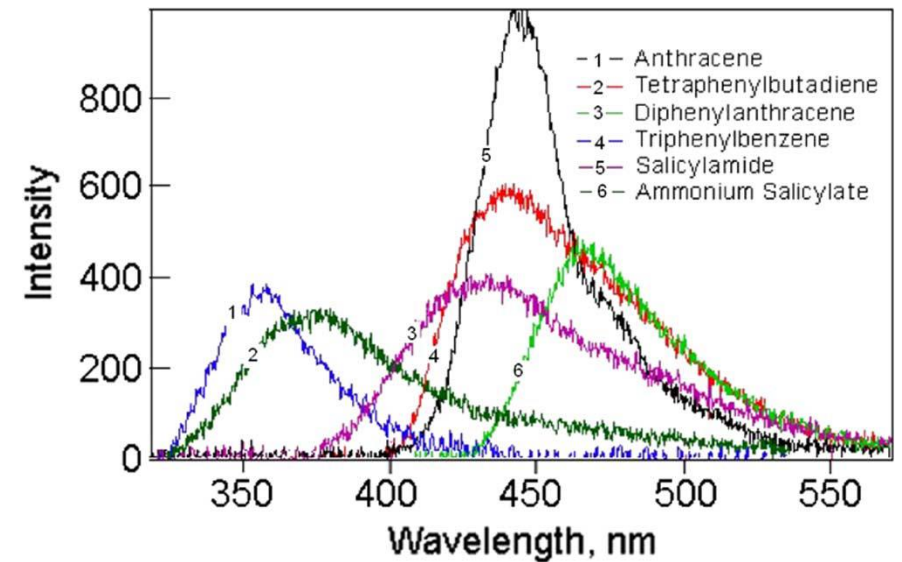


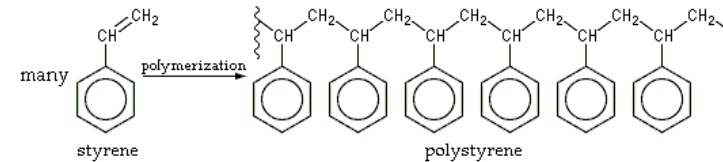
Fig. 4. Neutron/Gamma separation profile for triphenylbenzene (top) and tetraphenylbutadiene (bottom).



# LIQUID AND PLASTIC SCINTILLATORS

- The bulk of the mass is a solvent which absorbs the ionization energy
  - Benzene (liquid), Linear alkylbenzene (liquid), polystyrene (plastic), PVT (plastic), etc.
- Non-radiative dipole-dipole transfer to primary fluor molecules
  - Another organic scintillator added (1%/Volume)
  - Foerster Resonant Energy Transfer (next slide)
- Emits a photon which is absorbed and re-emitted by the wavelength shifter
  - Yet another organic scintillator (0.1%)

- Example: polystyrene



- The radiative and non-radiative (quenching) transfer rates for polystyrene,  $k_R$  and  $k_{NR}$ :

$$k_R = 1.98 \times 10^6 s^{-1}$$

$$k_{NR} = 5.02 \times 10^7 s^{-1}$$

- The radiative yield quantum efficiency,  $q_R$ , and decay time,  $\tau_R$

$$q_R = \frac{k_R}{k_R + k_{NR}} = 0.038$$

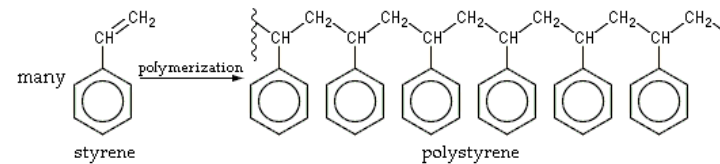
$$\tau_R = \frac{1}{k_R} = 500 ns$$

Terrible!

# LIQUID AND PLASTIC SCINTILLATORS

- The bulk of the mass is a solvent which absorbs the ionization energy
  - Benzene (liquid), Linear alkylbenzene (liquid), polystyrene (plastic), PVT (plastic), etc.
- Non-radiative dipole-dipole transfer to primary fluor molecules
  - Another organic scintillator added (2%/Volume)
  - Foerster Resonant Energy Transfer (next slide)
- Emits a photon which is absorbed and re-emitted by the wavelength shifter
  - Yet another organic scintillator (0.1%)

- Example: polystyrene



- The radiative and non-radiative (quenching) transfer rates for polystyrene,  $k_R$  and  $k_{NR}$ :

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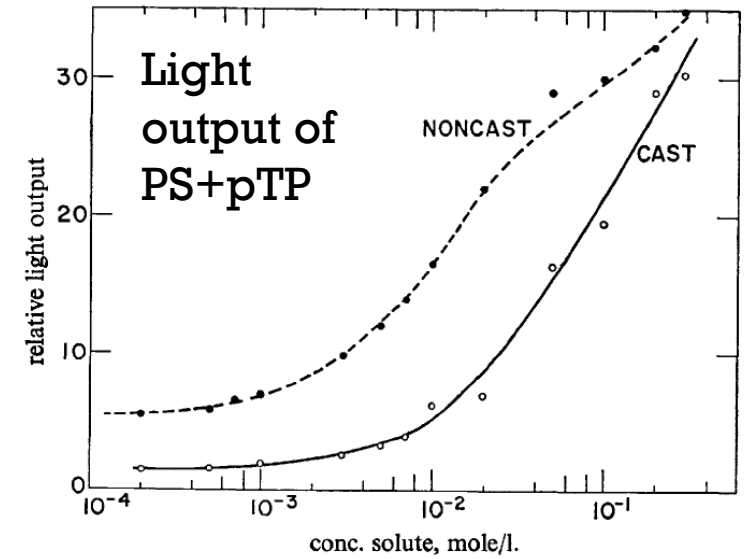
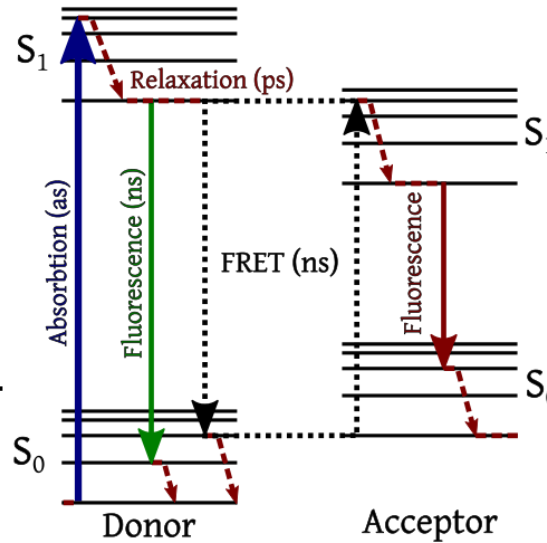
$$q_R = \frac{k_R}{k_R + k_{NR}} = 0.038$$

$$\tau_R = \frac{1}{k_R} = 500 \text{ ns}$$

Terrible!

# FRET

- can add a second organic scintillator to compete for the energy transfer via **Förster Resonance Energy Transfer (FRET)**
- Non-radiative dipole–dipole coupling between donor and acceptor molecule
  - Like near-field communication
  - Exchange of a virtual photon
- Transfer rate and efficiency is concentration dependent (maximize)
  - Beware of self-absorption



1.—Light output against concentration of p-terphenyl in polystyrene.

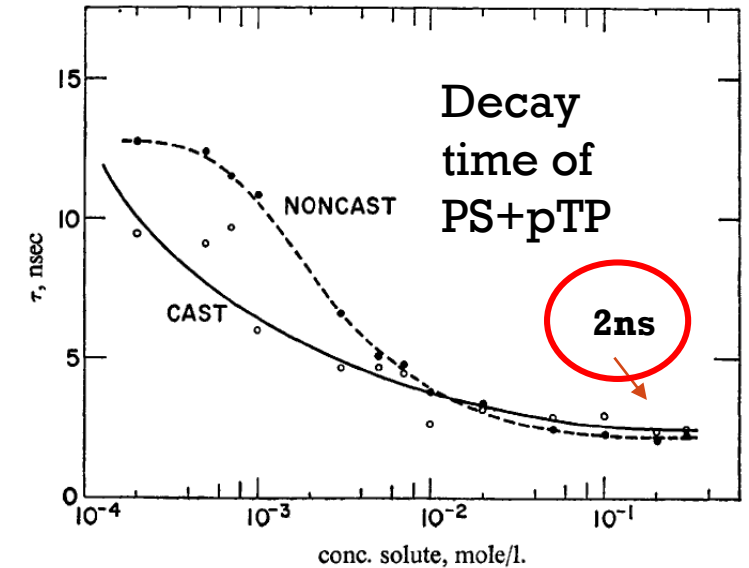
$$c \approx 10^{-3} \rightarrow 10^{-1} \text{ mol/l}$$

$$k_T \approx 10^9 \rightarrow 10^{10} \text{ l mol}^{-1} \text{ sec}^{-1}$$

$$q_T = \frac{k_{TC}}{k_R + k_{NR} + k_{TC}} \approx 1$$

**All the energy is transferred to the second fluor and very quickly (<ns)!**

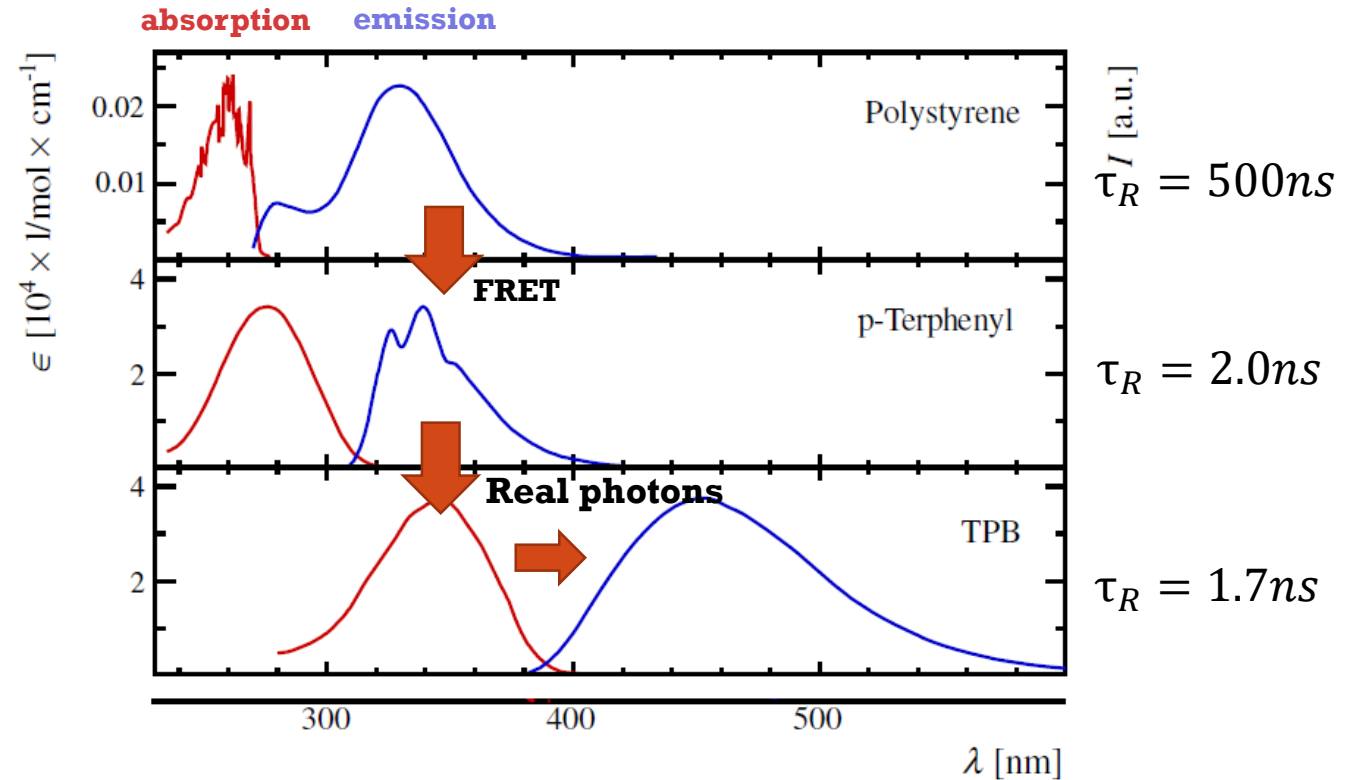
$$\tau_{PS} = \frac{1}{k_R + k_{NR} + k_{TC}}$$



Louis J. Basile  
Trans. Faraday Soc., 1964,60, 1702-1714

# LIQUID AND PLASTIC SCINTILLATORS

- The bulk of the mass is a solvent which absorbs the ionization energy
  - Benzene (liquid), Linear alkylbenzene (liquid), polystyrene (plastic), PVT (plastic), etc.
- Non-radiative dipole-dipole transfer to primary fluor molecules
  - Another organic scintillator added (2%/Volume)
  - Foerster Resonant Energy Transfer (next slide)
- Emits a real photon which is absorbed and re-emitted by the wavelength shifter
  - Yet another organic scintillator (0.1%)
  - Longer wavelength = better transmission



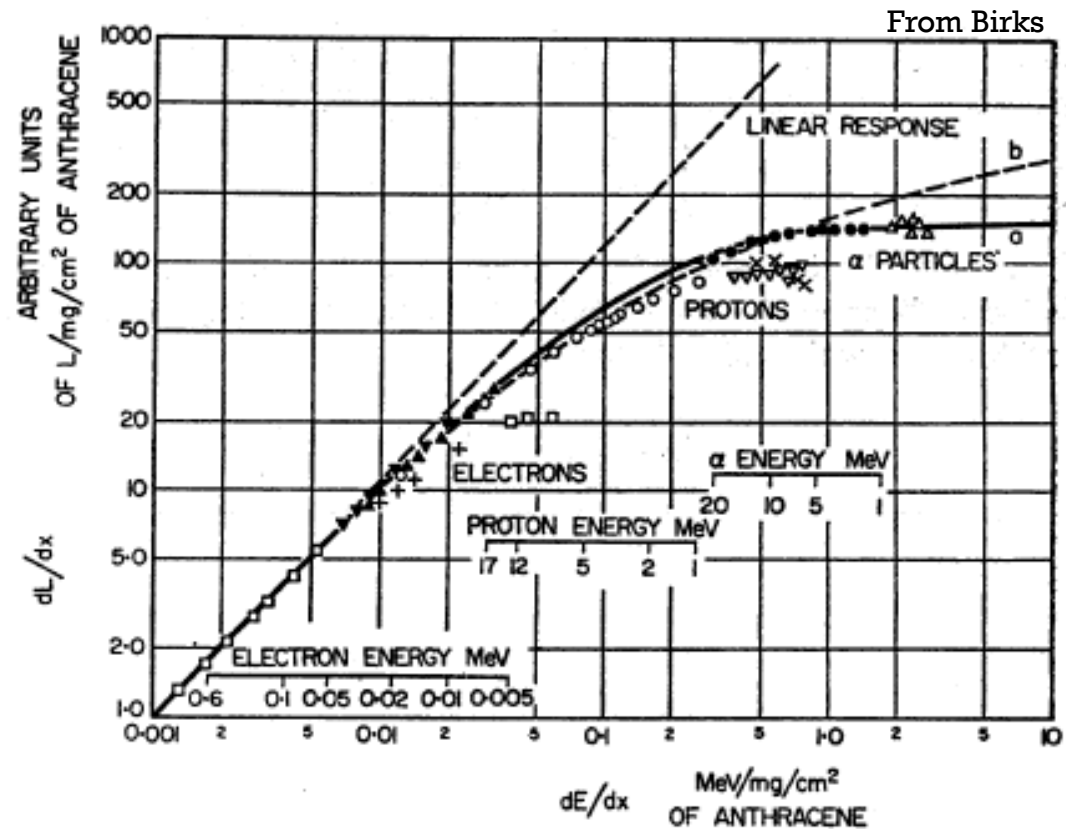
$$\tau_{total} = 2.0 \oplus 1.7 ns = 2.8ns$$



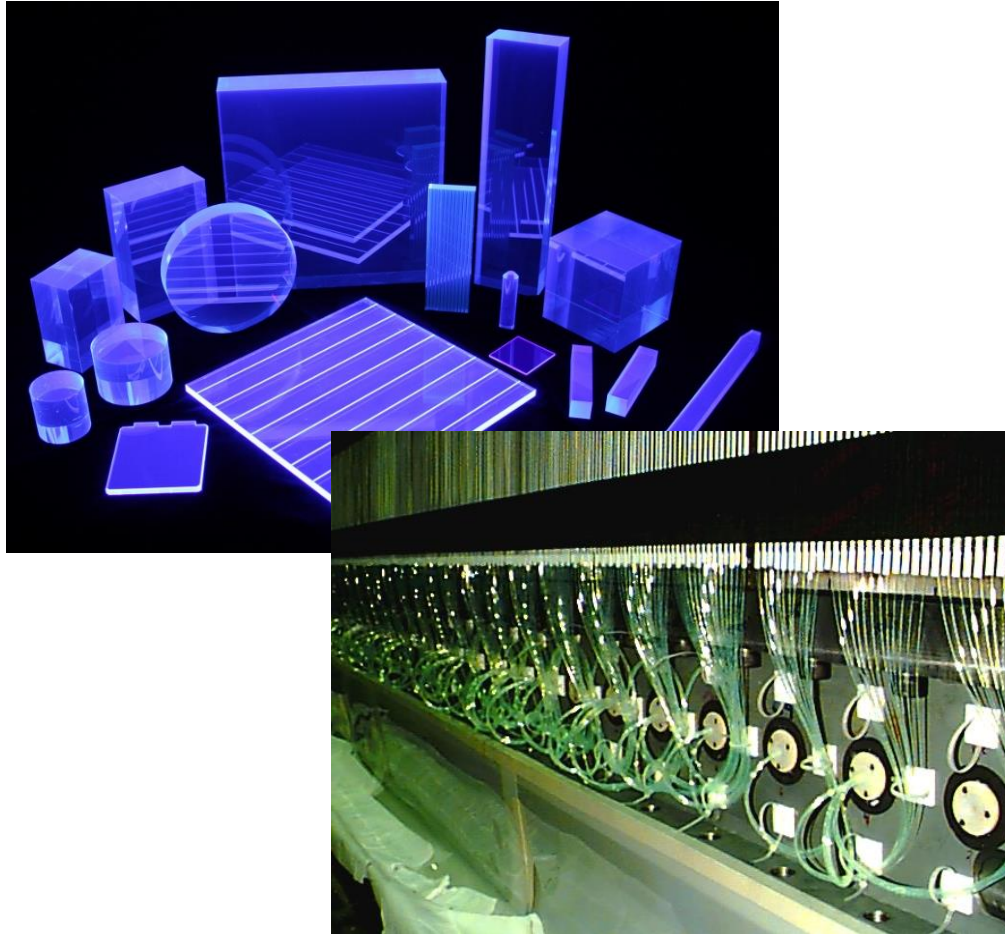
# SATURATION: BIRKS' LAW

- Assume linear light output for small  $dE/dx$ 
  - $dL/dx = A dE/dx$
- density of excitation is proportional to energy density
  - $B dE/dx$
- $k$  is the fraction of excitation that is quenched
  - Interaction between nearby excitations results in non-radiative transfers

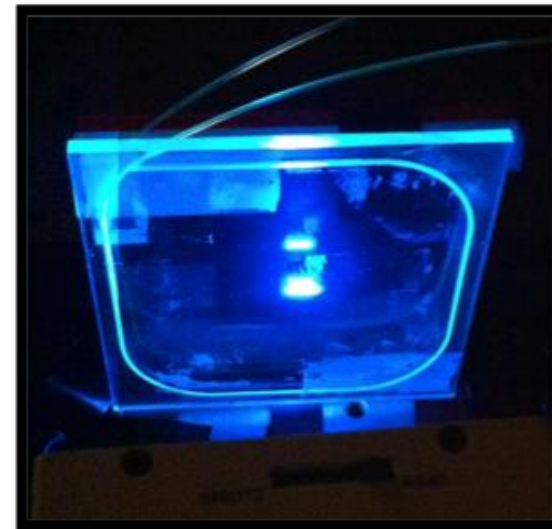
$$\frac{dL}{dx} = \frac{A \frac{dE}{dx}}{1 + kB \frac{dE}{dx}}$$



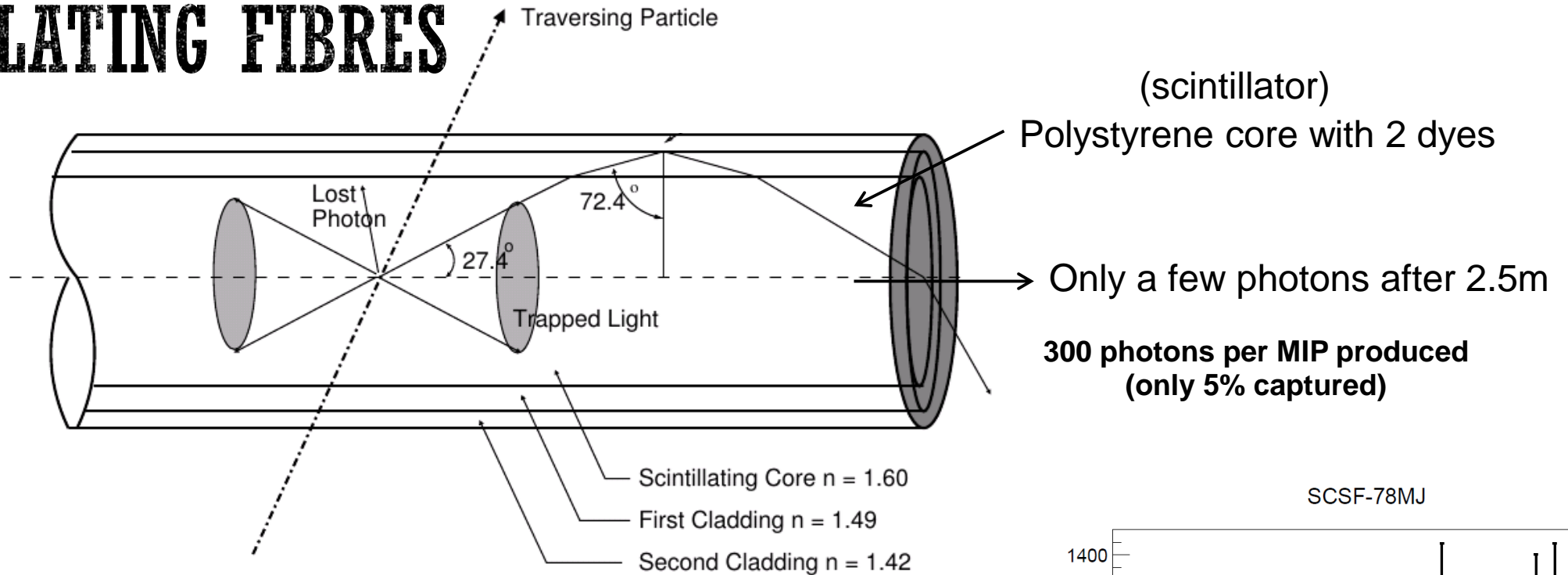
# FORMS



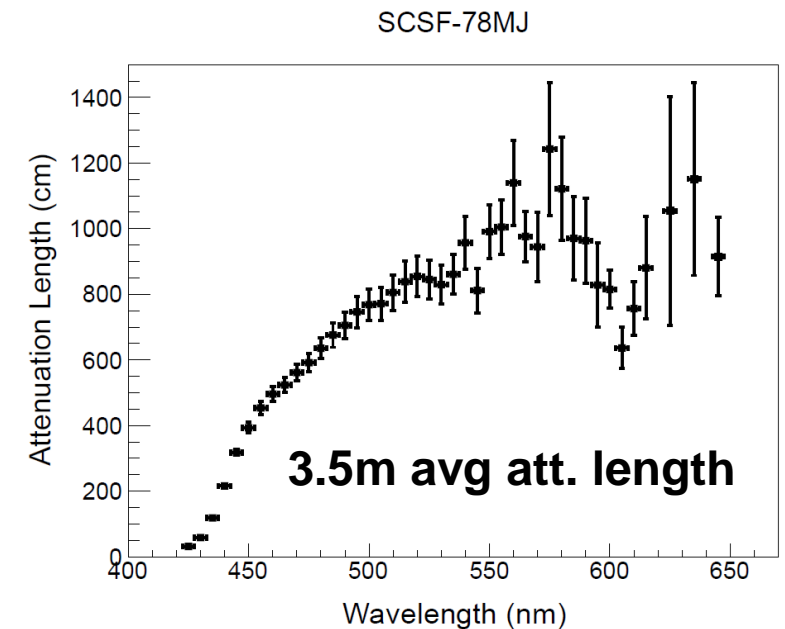
- Fast and cheap (relatively)
- Simpler to produce compared to crystals
  - Cast, injection mold or extrusion is possible
  - Large sheets or volumes can be produced relatively quickly
- Standard operating practices:
  - couple a waveguide and a PMT
  - Embed wavelength shifting fibres and transport to smaller size photodetectors



# SCINTILLATING FIBRES

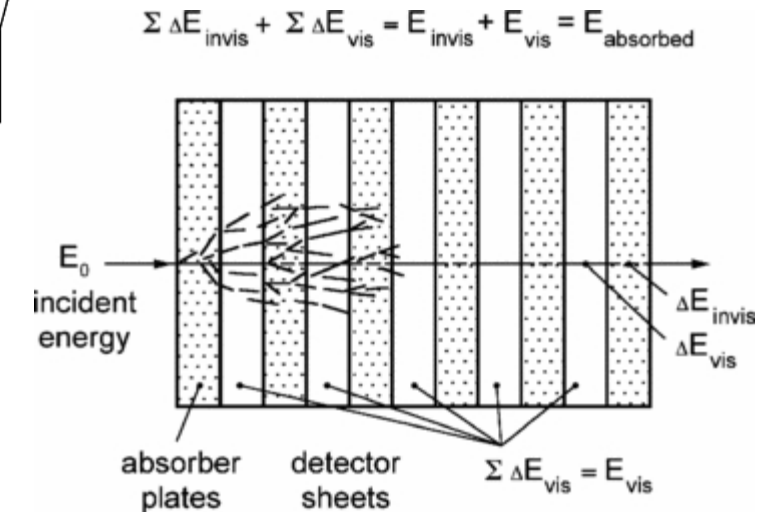
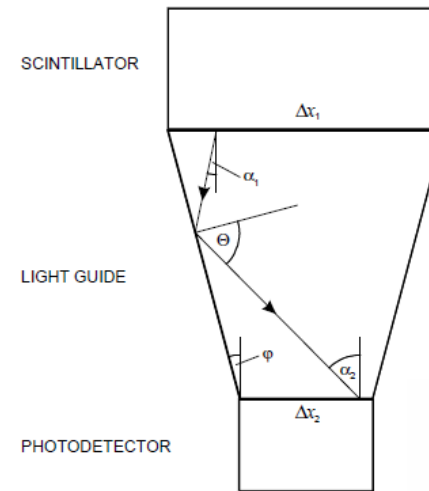


- Scintillator is melted and extruded with a small diameter (0.25 – 2mm)
  - Round or square profiles
- One or two lower refractive index plastics added to improve collection efficiency (3-5% eff.)



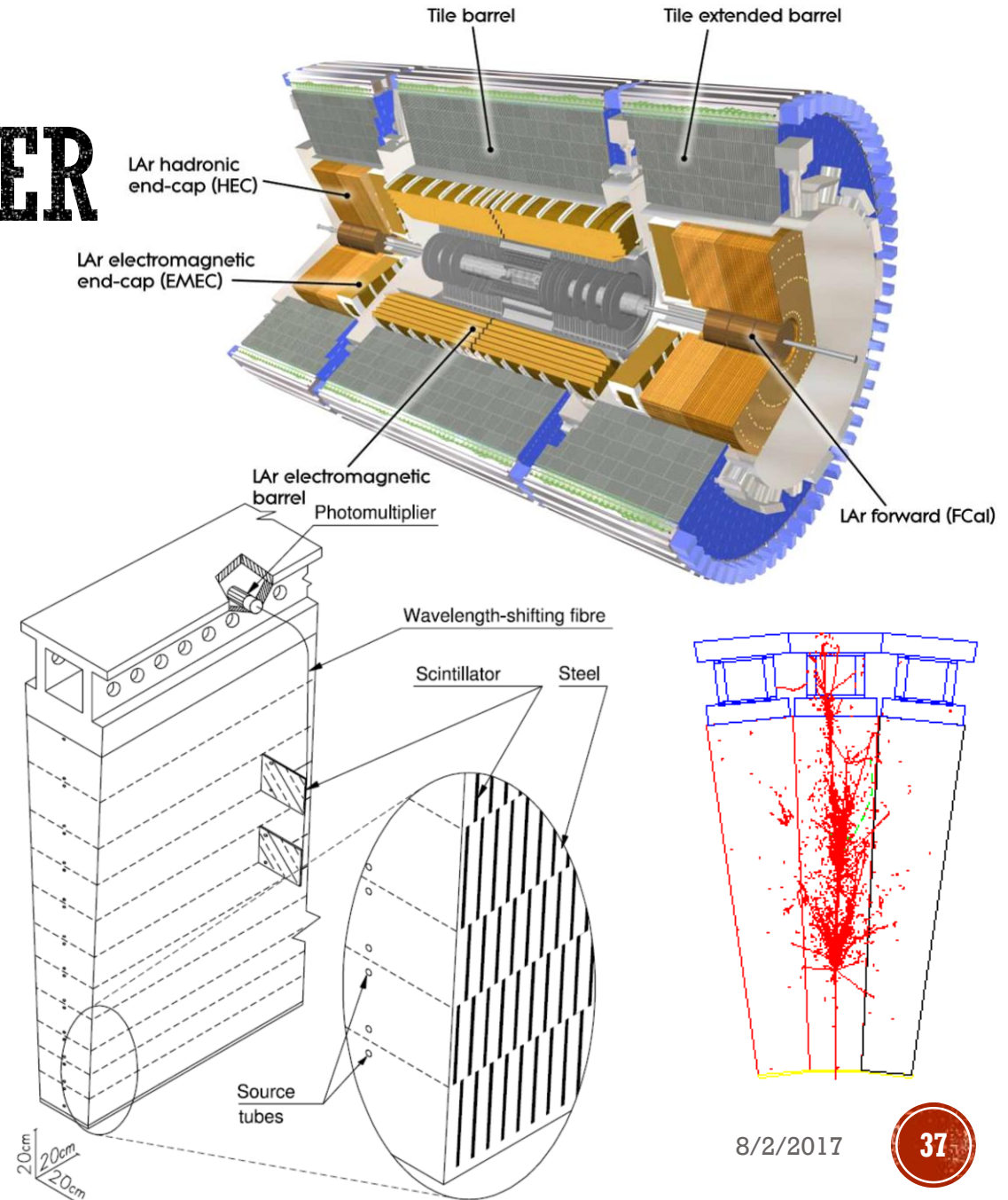
# EXAMPLE DETECTORS

- Triggers for timing
  - Low Z, density, fast decay time
- Sampling Calorimeters
  - Layers of scintillator + high-Z passive material
    - Tile calorimeters
    - Spaghetti calorimeters (fibres)
- Neutron detectors (liquid, some new plastic)
  - Good hydrogen content
- Liquid Scintillator Baths
  - Easy to add additional materials
  - Neutrino detector (SNO+)



# ATLAS TILE CALORIMETER

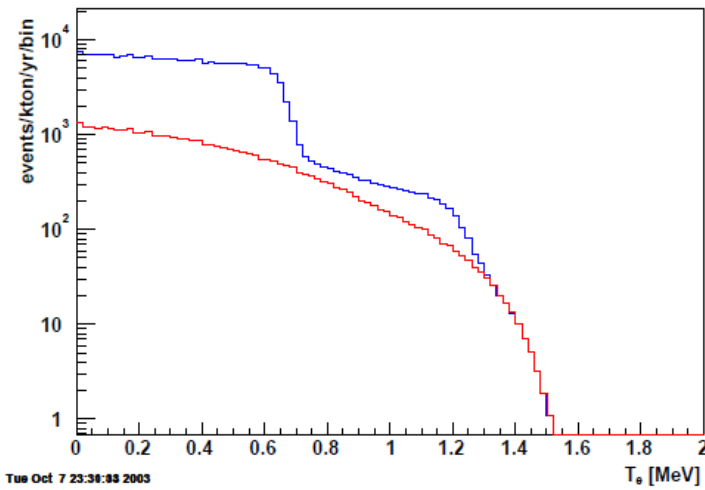
- Steel + scintillator + w.s. fibres
- Hadronic calorimeter
- $\frac{\delta E}{E} \sim \frac{100\%}{\sqrt{E}} \oplus 10\%$  Testbeam, ATLAS NIMA 621, Sep 2010, 134-150
- 460,000 tiles in 5182 cells (60 tons)
- Polystyrene + 1.5% PTP and 0.04% POPOP
  - Injection molding
  - <http://cds.cern.ch/record/1075711/files/cer-002731189.pdf>
- PMT as photodetector



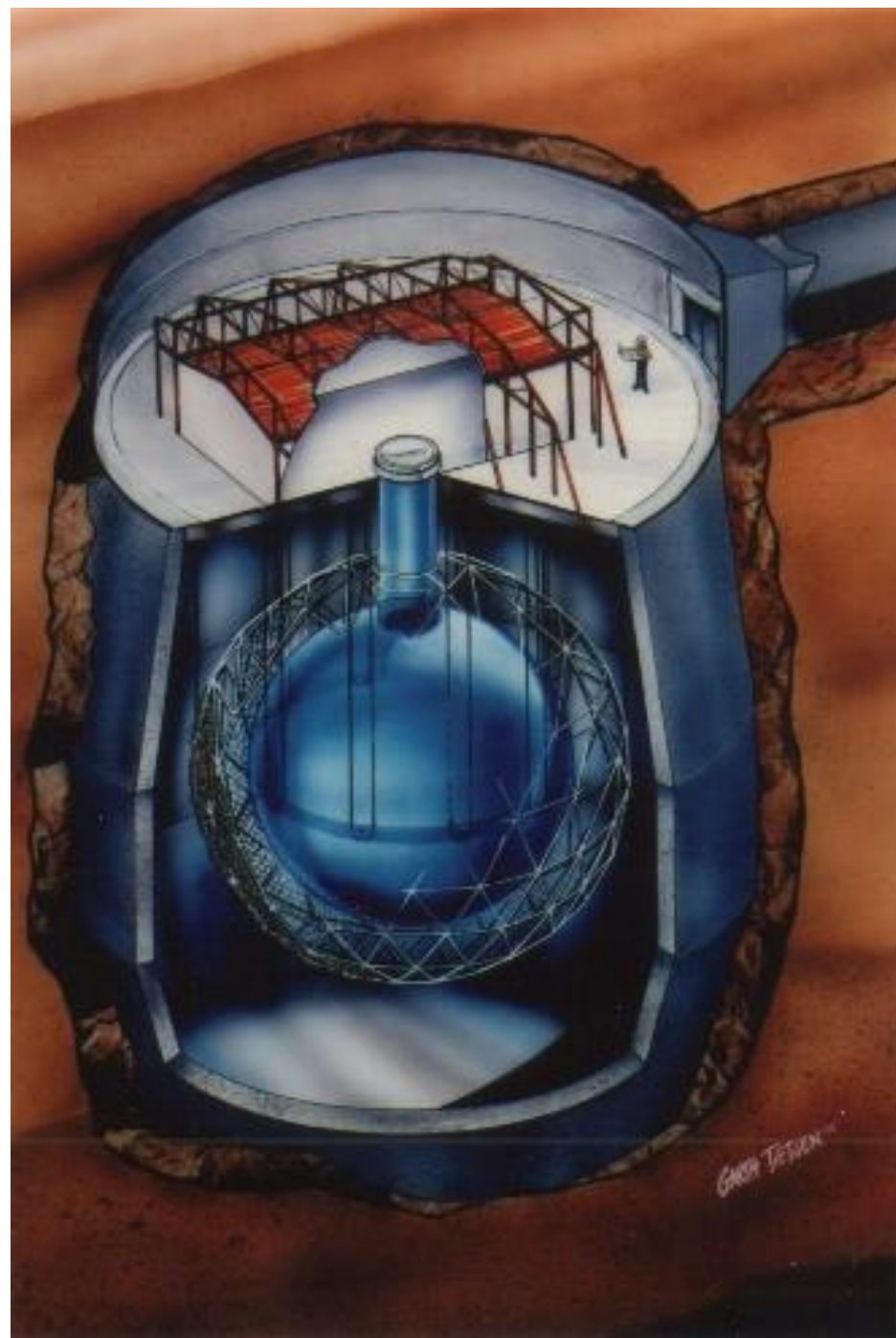
# SNO+

- 1 kiloton of liquid scintillator (linearalkylbenzene-LAB)
- 50x more light than Cherenkov
- Solar-, geo-neutrino study and neutrinoless double beta decay search (SNO++)

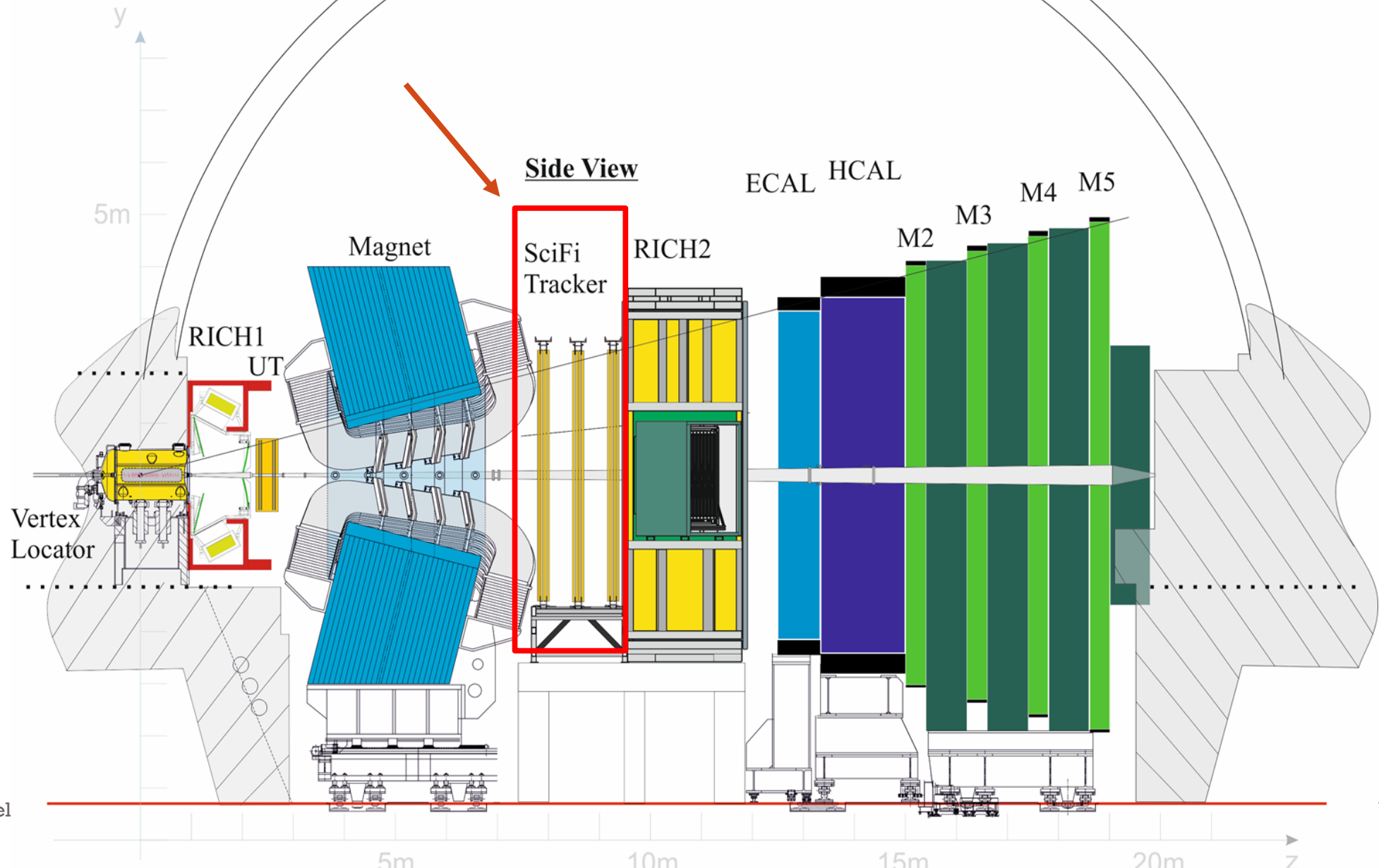
<sup>7</sup>Be, pep and CNO Recoil Electron Spectrum



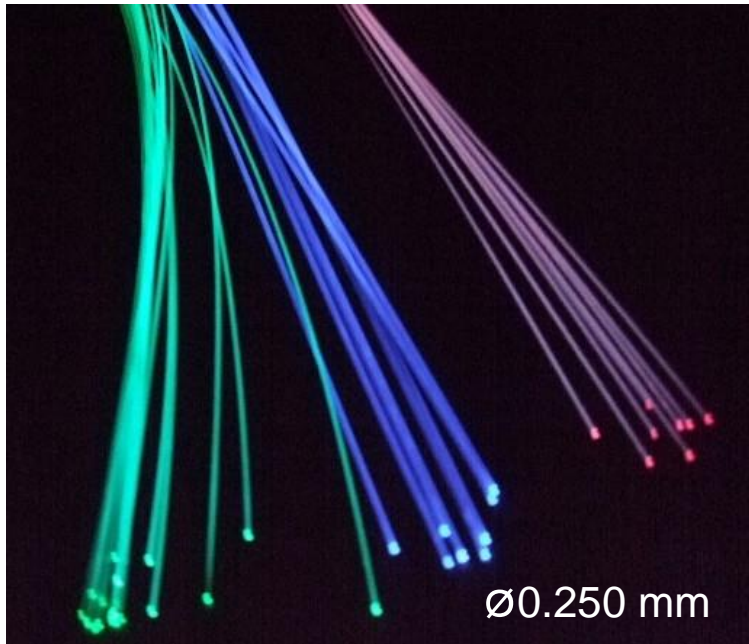
Chen, Nuclear Physics B (Proc. Suppl.) 145 (2005) 65–68



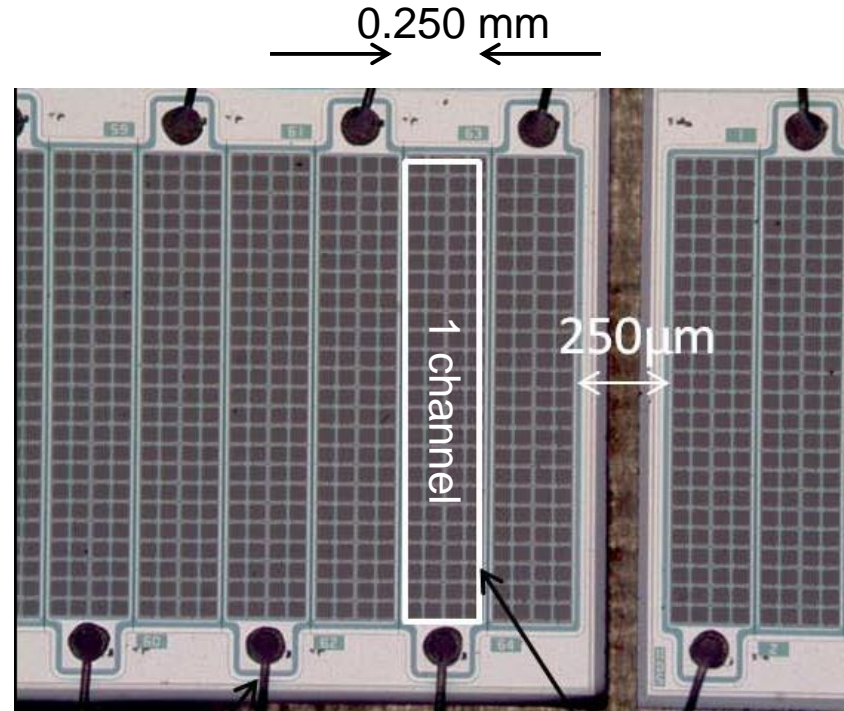
# LHCb Upgraded Spectrometer



# THE SCIFI TRACKER



Scintillating fibres



An array of pixelated silicon photomultipliers

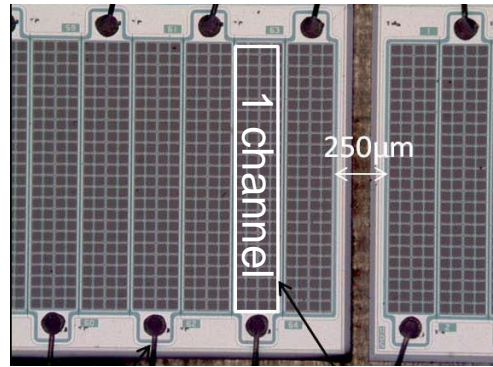


# TRACKER REQUIREMENTS

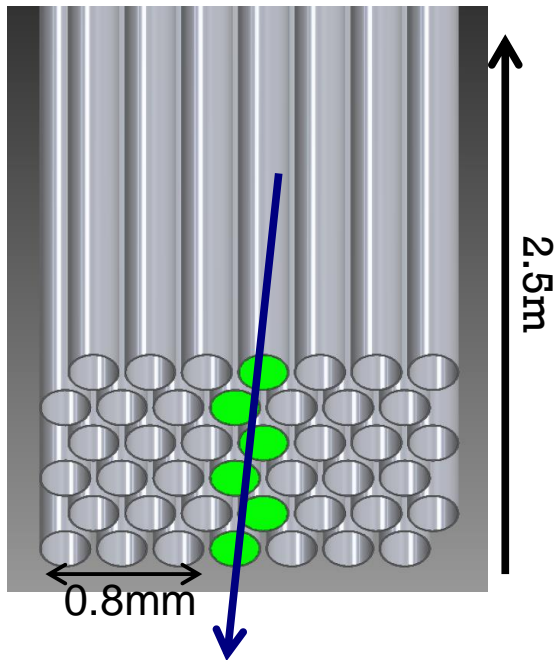
- Hit detection efficiency: single hits ~ 99% → Track reconstruction
- Low material budget for single detector layer ~1% X0 → Multiple scattering  
Nuclear interactions
- Spatial resolution better: 100 μm in x-direction → Momentum resolution
- 40 MHz readout without dead time → Data rate / efficiency
- Radiation environment: Fibres: up to 35 kGy, → Signal/noise ratio
- SiPMs: approx.  $1 \cdot 10^{12}$  n/cm<sup>2</sup> fluence + 100 Gy ionizing dose

# BASIC PRINCIPLE

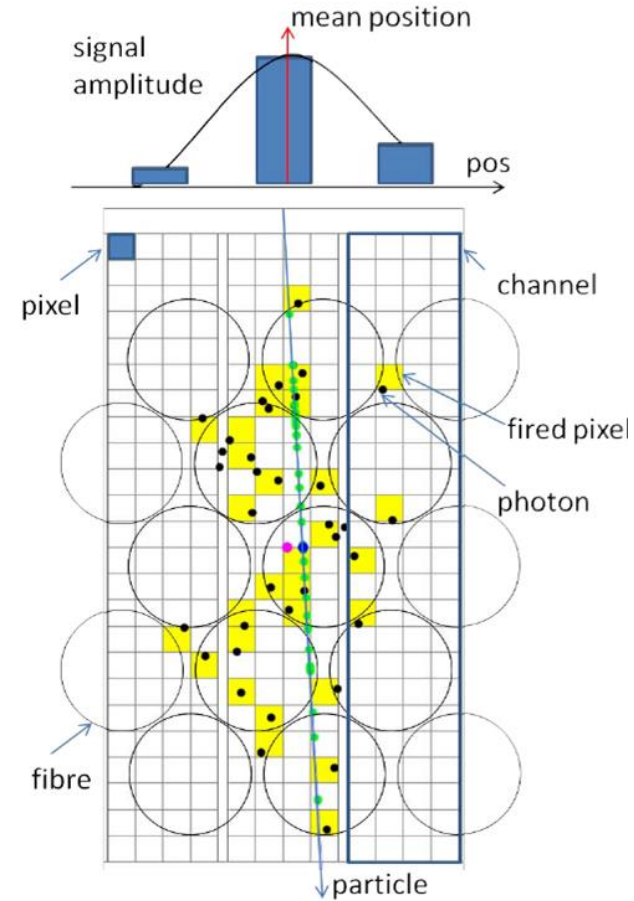
SiPM array



Scintillating Fibres  
(0.250mm diameter)

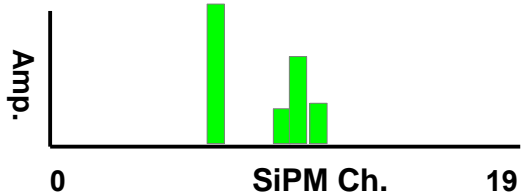


Signal cluster

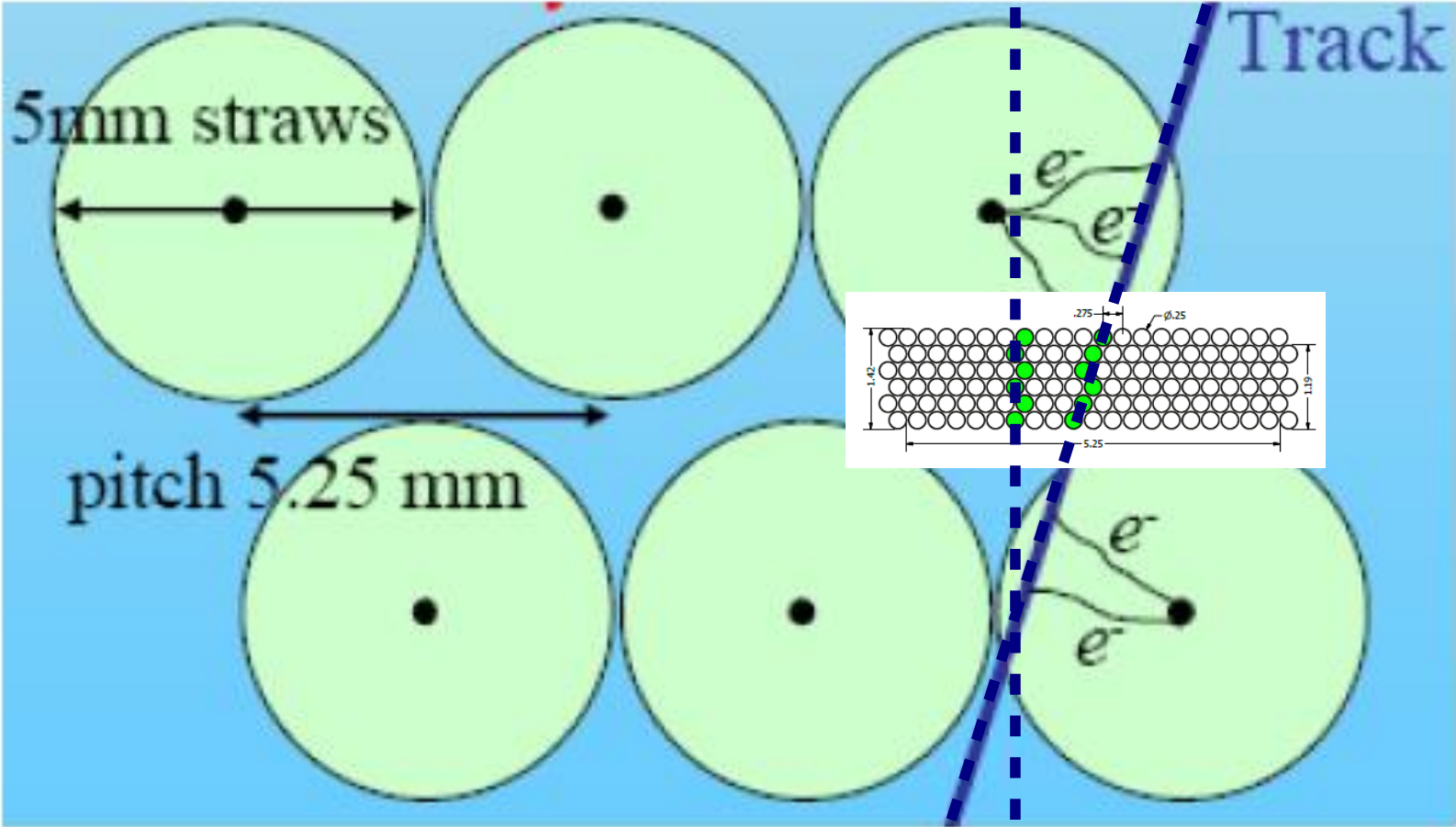


Typically one observe 15-20 photoelectrons for 5 layers of fibre

Outer Tracker straws



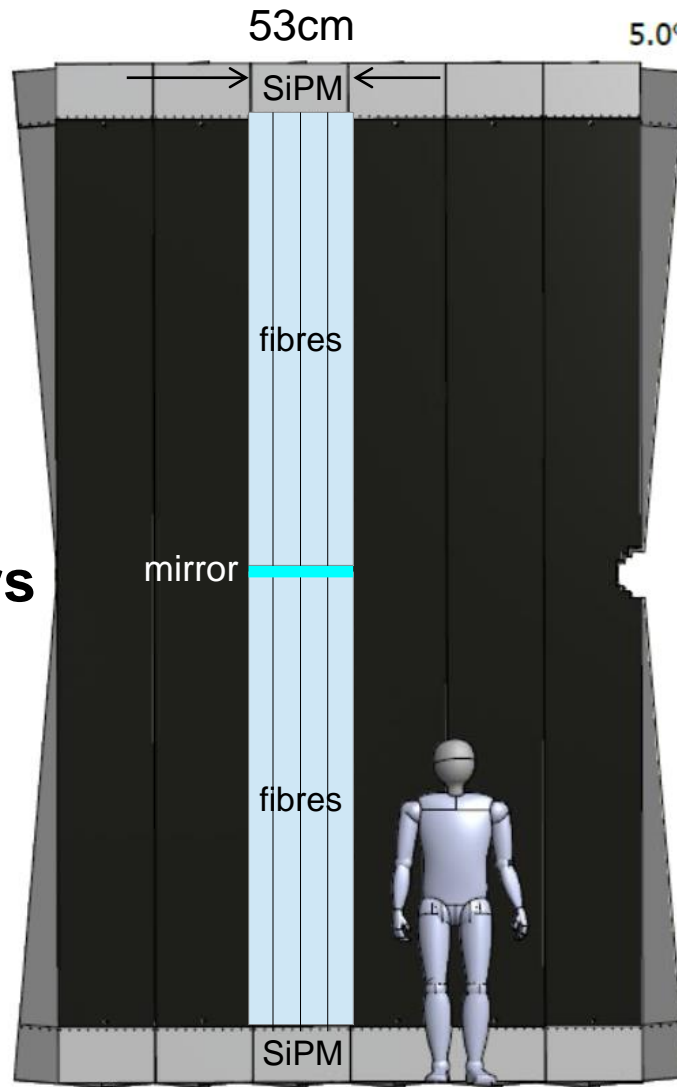
127



Approximately to relative scale

XUVX

x 3 = 12 layers



5.0°

Modules are assembled in Heidelberg



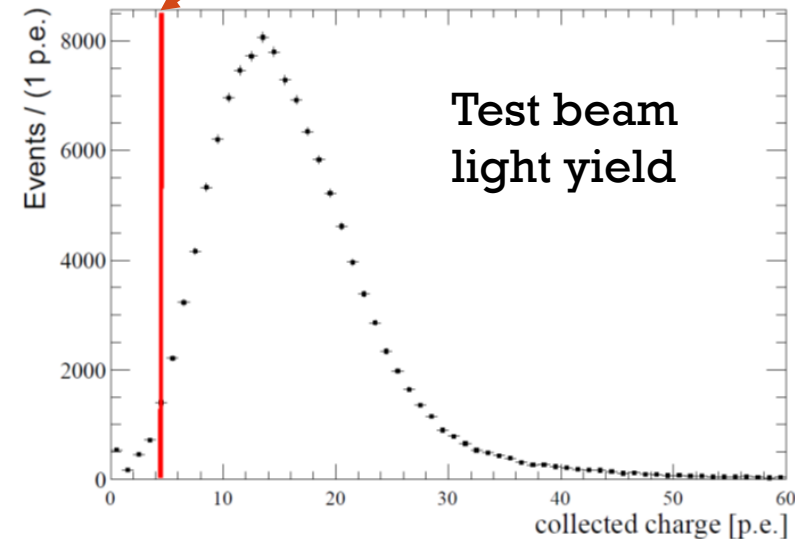
# CHALLENGES

## ▪ Timing

- 2.8ns decay time of scintillator
- 2.5m path length (15ns spread in arrival times)
- 10ns pre-amp shaping time
  - Some charge is collected in the neighbouring bunch crossing
- 25ns bunch crossing
  
- PACIFIC asic being designed in Heidelberg

## ▪ Light yield / hit efficiency

- 16-18 mean photoelectrons detected near the mirror
- $\sqrt{N}$  fluctuation in light yield
- Threshold for dark noise suppression



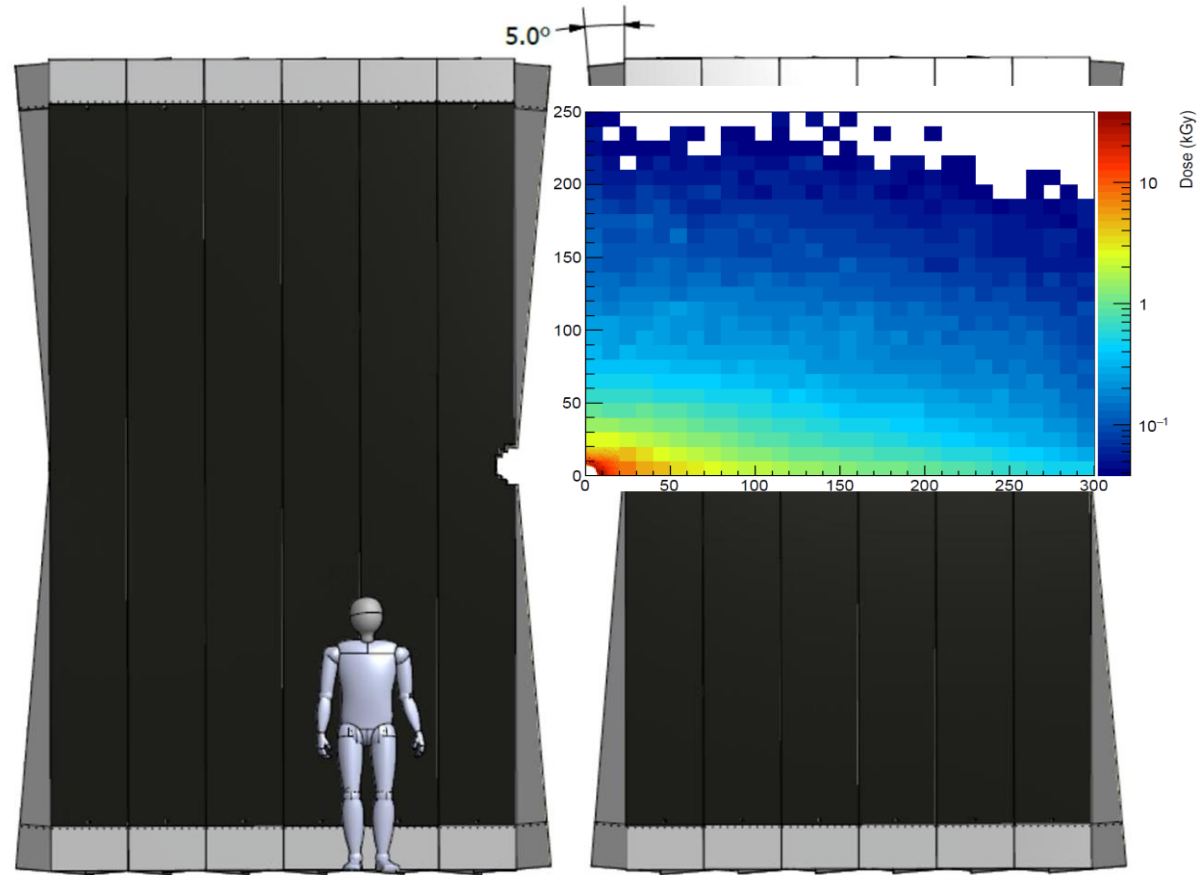
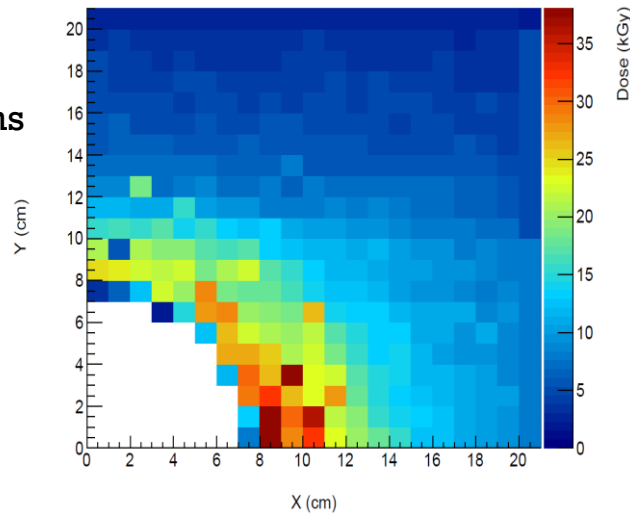
# IONIZING RADIATION

- Highest ionizing radiation damage is near the beam pipe

Max = 35 kGy @ 50fb<sup>-1</sup>

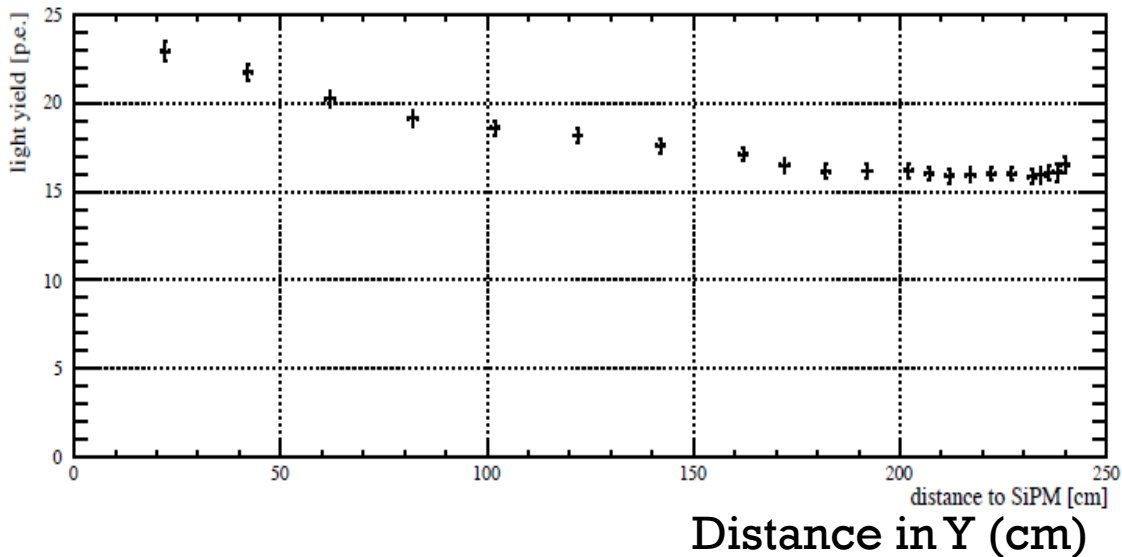
- Reduces quickly away from the beampipe

FLUKA simulations

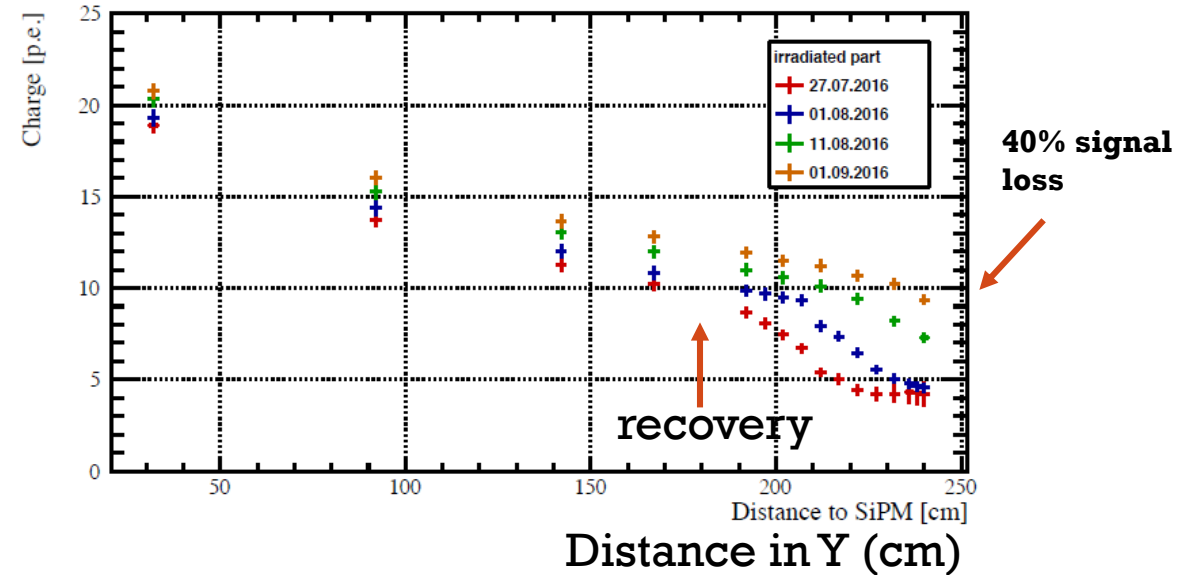


# RADIATION DAMAGE

- Before irradiation



- After 50 fb<sup>-1</sup> irradiation (in worst region)
  - Transmission decreases from additional absorption centers



Improvements →

- New SiPM with 50% PDE (vs 40%)
- Production fibre light yield has increased 10%

# CONCLUSION

- Scintillators play a large roll in radiation detection and High Energy physics
- The physics is based on solid state crystals (inorganic) or organic molecules
- The scintillator is chosen based on the application
  - A wide range of properties
- The LHCb Scintillating Fibre tracker is the largest fibre tracker ever built
  - Must operate in a demanding environment = many challenges



# GLOSSARY

- An **exciton** is a bound state of an electron and an electron hole which are attracted to each other by the electrostatic Coulomb force. It is an electrically neutral quasiparticle that exists in insulators, semiconductors and in some liquids. The exciton is regarded as an elementary excitation of condensed matter that can transport energy without transporting net electric charge. - Wikipedia
- a **phonon** is a collective excitation in a periodic, elastic arrangement of atoms or molecules in condensed matter, like solids and some liquids. Often designated a quasiparticle, it represents an excited state in the quantum mechanical quantization of the modes of vibrations of elastic structures of interacting particles. – Wikipedia
- **Hygroscopy** is the phenomenon of attracting and holding water molecules from the surrounding environment, which is usually at normal or room temperature. This is achieved through either absorption or adsorption with the absorbing or adsorbing substance becoming physically changed somewhat. This could be an increase in volume, boiling point, viscosity, or other physical characteristic or property of the substance, as water molecules can become suspended between the substance's molecules in the process. --Wikipedia