# Gamma-ray spectroscopy

Alison Bruce University of Brighton

Aim is to give an overview for non-specialists but also some details that specialists might find useful.

### Contents:

- Basic gamma-ray properties, observables
- Methods of producing the nuclei of interest (not an exhaustive list)
- Gamma-ray interactions in matter
- Detector types
- Detector arrays
- Measurement techniques:

Angular correlation, angular distribution

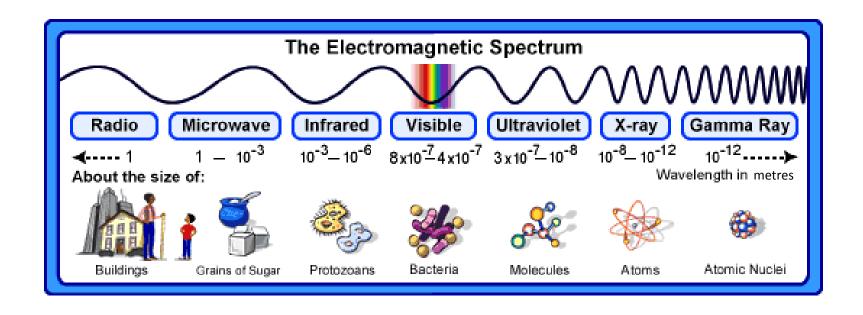
Linear polarisation

Lifetime measurements: Doppler Shift Attenuation Method

Recoil Distance Method

Electronic timing

# What is a gamma ray?

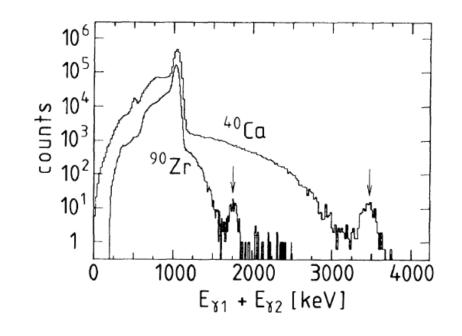


Gamma ray: high frequency / short wavelength electromagnetic radiation.

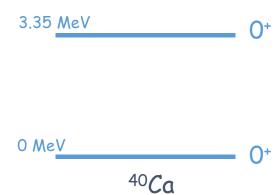
Useful as a probe of the nucleus as the electromagnetic interaction is well understood and only weakly perturbs the nucleus.

# Gamma rays carry spin 1

which leads to interesting cases where a 0<sup>+</sup> state is the lowest excited level:



Sum-energy spectrum in coincidence with protons populating the states of interest.



Schirmer et al., PRL 53 (1984) 1897 See also  $^{137}$ Ba, Pietrella et al., Nature 526 (2015) 406

# What is gamma-ray spectroscopy?

We study gamma rays emitted from excited nuclei to obtain information about:

Transition energies and coincidence relationships Level structure

Transition rates
Lifetimes, quadrupole moment

Angular correlations and linear polarisations Spin, Parity (also magnetic moments).

Transition branching ratios, mixing ratios
Wavefunctions, transition matrix elements, etc

# Why study gamma rays?

Gamma rays provide a superb probe for nuclear structure!

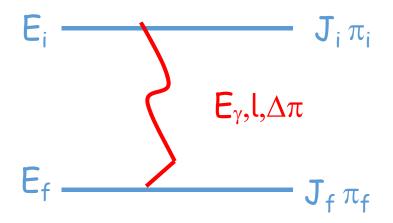
Relatively easy to detect with good efficiency and resolution.

Emitted by almost all low-lying states.

Penetrating enough to escape from target chambers to reach detectors.

Gamma rays arise from EM interactions and allow a probe of structure without large perturbations of the nucleus.

No model dependence in the interaction (EM is well understood).



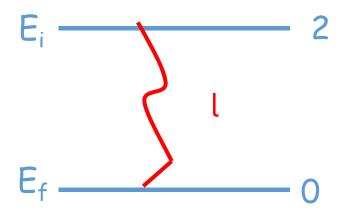
The energy of the gamma ray is given by  $E_i$ - $E_f$ 

The angular momentum carried away is given by  $J_i$ - $J_f$ 

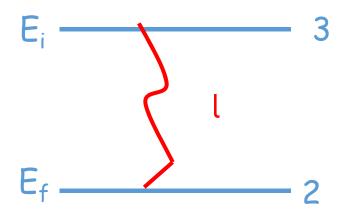
From conservation of angular momentum:

$$|J_i - J_f| \le \ell \le J_i + J_f$$

where  $\ell$  is the multipolarity of the transition

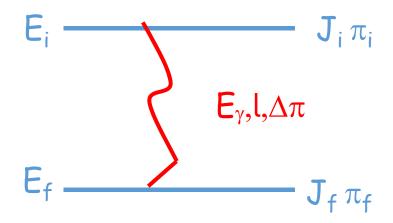


Here  $\Delta J = 2$  and  $\ell = 2$  so we say this is a stretched transition



Here 
$$\Delta J = 1$$
 but  $\ell = 1,2,3,4,5$ 

and the transition can be a mix of 5 multipolarities

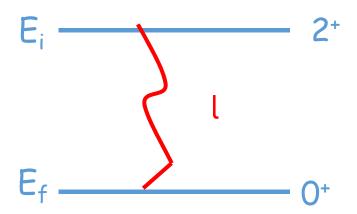


### Electromagnetic transitions:

$$\Delta\pi$$
 (electric) =  $(-1)^{\ell}$ 

$$\Delta\pi$$
 (magnetic) =  $(-1)^{\ell+1}$ 

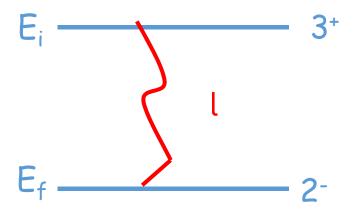
$\Delta\pi$		E1	M2	<b>E3</b>	M4
	NO	M1	<b>E2</b>	M3	<b>E4</b>



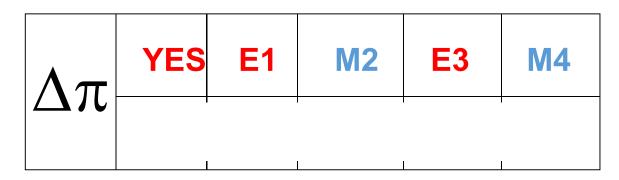
 $\ell$  = 2 and no change in parity

$\Delta\pi$				
	M1	<b>E2</b>	M3	<b>E4</b>

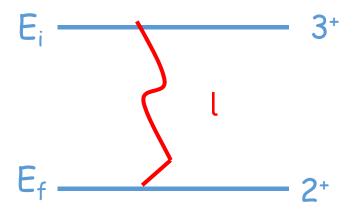
pure (stretched) E2



Here  $\Delta J = 1$  but  $\ell = 1,2,3,4,5$ 



Mixed E1/M2/E3/M4/E5



Here  $\Delta J = 1$  but  $\ell = 1,2,3,4,5$ 

$\Delta\pi$	I	1	1	
	M1	<b>E2</b>	M3	<b>E4</b>

Mixed M1/E2/M3/E4/M5

3+ -> 2+: mixed M1/E2/M3/E4/M5

3+ -> 2-: mixed E1/M2/E3/M4/E5

In general only the lowest 2 multipoles compete

and (for reasons we will see later)

l+1 multipole generally only competes if it is electric so:

3+ -> 2+: mixed M1/E2

3+ -> 2-: almost pure E1 (v. little M2 admixture)

Amount of mixing is given by the mixing ratio  $\delta$ 

given by 
$$\delta = \frac{\left\langle \psi_f \| M(l+1) \| \psi_i \right\rangle}{\left\langle \psi_f \| M(l) \| \psi_i \right\rangle}$$

3+ -> 2+: M1/E2 admixture

%E2 = 
$$\frac{\delta^2}{1+\delta^2}$$

%M1 = 
$$\frac{1}{1+\delta^2}$$

# Not quite so basic:

Transition rate  $T[s^{-1}]$  given by:

$$T_{f\to i}(\sigma\lambda) = \frac{2(l+1)}{\varepsilon_0 l[(2l+1)!!]^2} \left(\frac{E_{\gamma}}{c}\right)^{2l+1} \langle \psi_f | M(\sigma\lambda) | \psi_i \rangle^2$$

Note the I dependence on the energy, the double factorial (what is that?)

$$\langle \psi_f | M(\sigma \lambda) | \psi_i \rangle^2$$

this is the bit that has the physics in it, it's the matrix element between the initial and final wavefunctions squared and is related to the  $B(\sigma\lambda)$  by

$$\mathsf{B}(\sigma\lambda) = \frac{1}{2J_i + 1} \sum_{M_i M_f} \langle \psi_f \big| M(\sigma\lambda) \big| \psi_i \rangle^2$$

# Not quite so basic:

Typical transition rates  $[s^{-1}]$ :

$$T(E1) = 1.59 \times 10^{15} (E_{\gamma})^3 B(E1)$$

$$T(E2) = 1.22 \times 10^9 (E_{\gamma})^5 B(E2)$$

$$T(M1) = 1.76 \times 10^{13} (E_y)^3 B(M1)$$

$$T(M2) = 1.35 \times 10^7 (E_y)^5 B(M2)$$

 $E_{\nu}$  in MeV

B(E $\lambda$ ) in e<sup>2</sup>fm<sup>2 $\lambda$ </sup>

B(M
$$\lambda$$
) in  $\left(\frac{e\hbar}{2Mc}\right)^2$  fm<sup>2 $\lambda$ -2</sup>

### Note:

- Electric transitions faster than magnetic
- Higher multipolarity => slower rate
- Transition probability proportional
   to transition energy => low-energy transitions
   are hard to observe and other processes
   e.g. internal conversion start to compete.

# Not quite so basic:

Single particle estimates (also called Weisskopf estimates)

$$T(E1) = 1.02 \times 10^{14} (E_{\gamma})^3 A^{2/3}$$

$$T(E2) = 7.26 \times 10^7 (E_y)^5 A^{4/3}$$

$$T(M1) = 3.18 \times 10^{13} (E_{\gamma})^3$$

$$T(M2) = 2.26 \times 10^7 (E_y)^5 A^{2/3}$$

Measuring level lifetimes gives us transition rates which can be interpreted as single particle (or collective) and hence give an indication of the type of motion

# Methods of producing the nuclei of interest

### Out-of-beam spectroscopy:

Nucleus is stopped Not many gamma rays emitted (Gamma-ray multiplicity low)

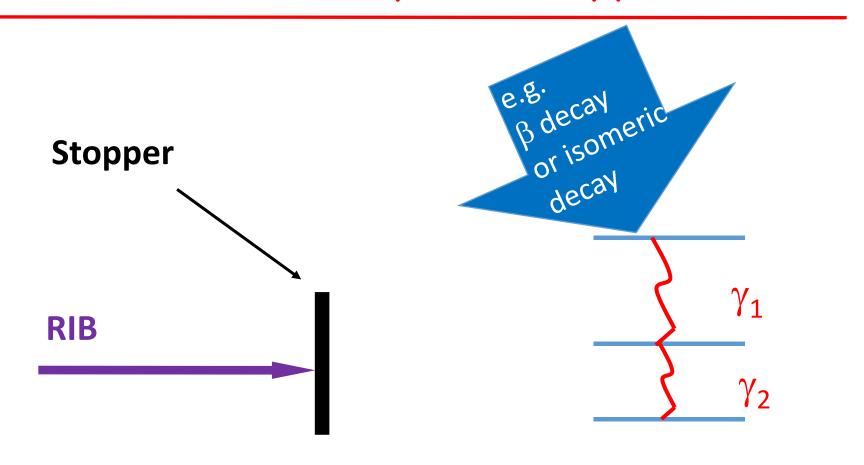
e.g. decay from a fission source, from stopped radioactive ion beams (ISOL or fragmentation), or de-excitation of isomeric states

### In-beam spectroscopy:

Nucleus is moving Lots of gamma rays emitted (Gamma-ray multiplicity high)

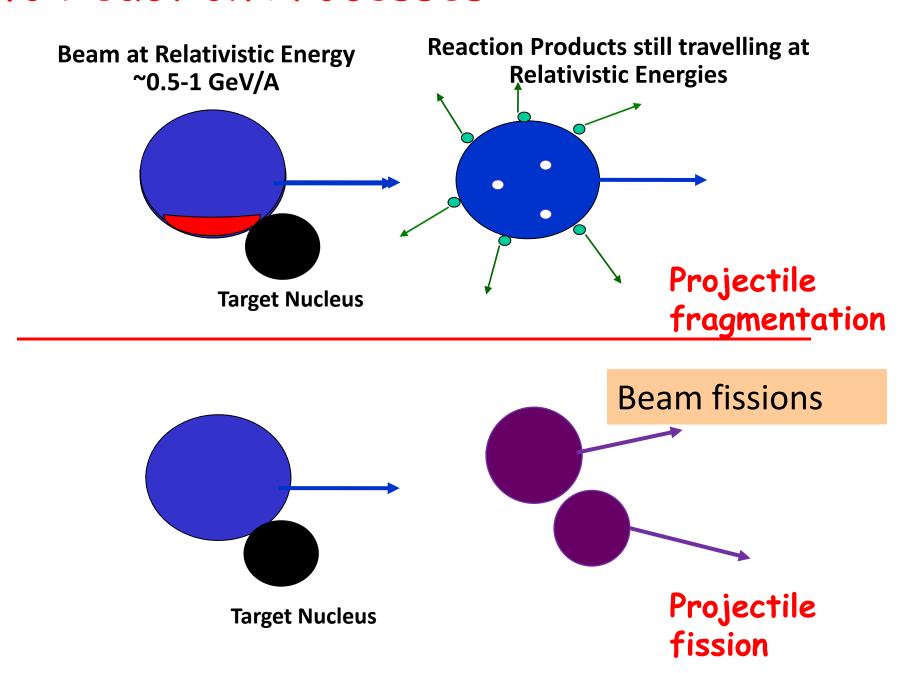
e.g. compound nucleus reaction, Coulomb excitation

# Out-of-beam spectroscopy

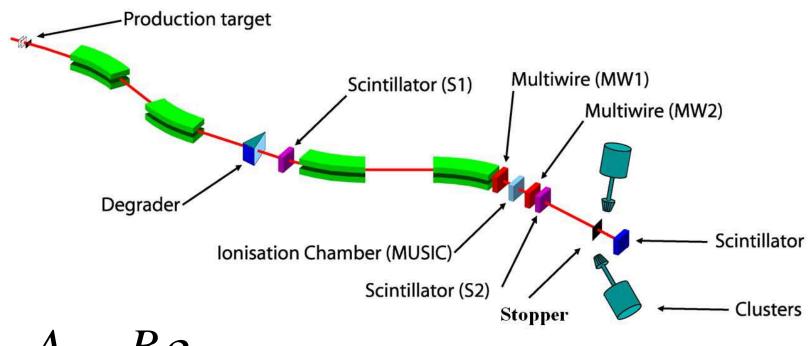


Get first information on lifetimes, decay modes, Q-values and scheme of excited levels

### Two Reaction Processes



# Ion-by-ion identification with e.g the FRS:



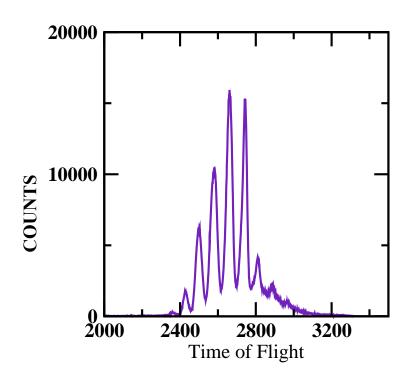
$$rac{A}{Z} \propto rac{B
ho}{eta\gamma}$$

FRS = Fragment Recoil Spectrometer at GSI

Cocktail of fragments Chemically independent

H.Geissel et al., NIM B70 (1992) 286.

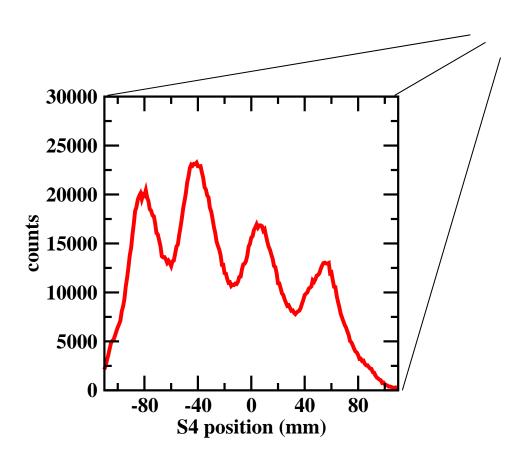
### Ion-by-ion identification with e.g the FRS:



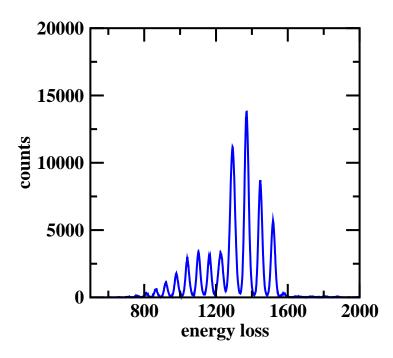
Time of flight

-> β

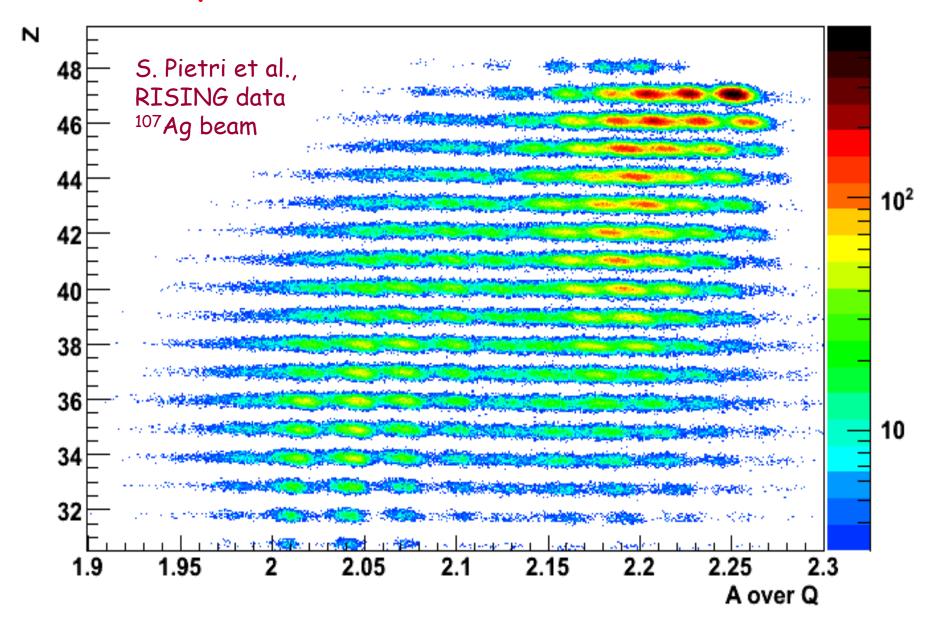
## Ion-by-ion identification with e.g the FRS:



# Ion-by-ion identification with e.g. the FRS:

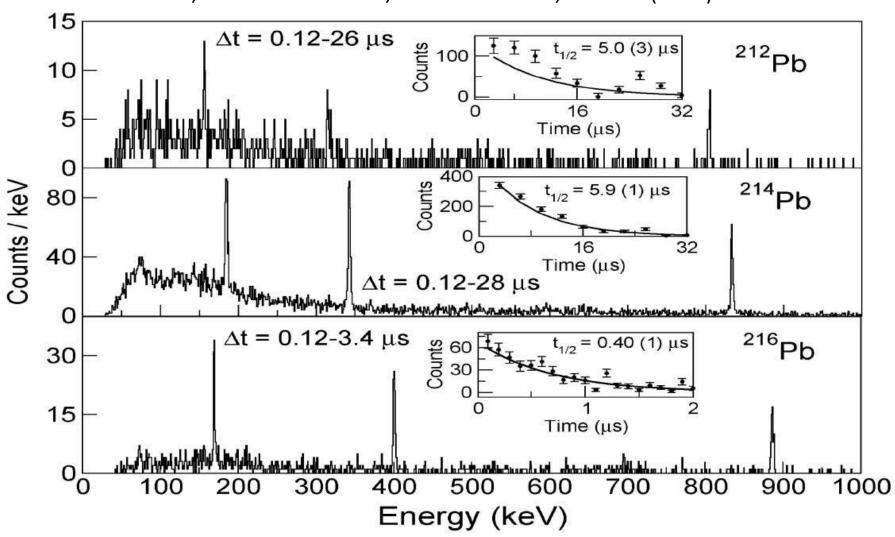


# Isotope identification



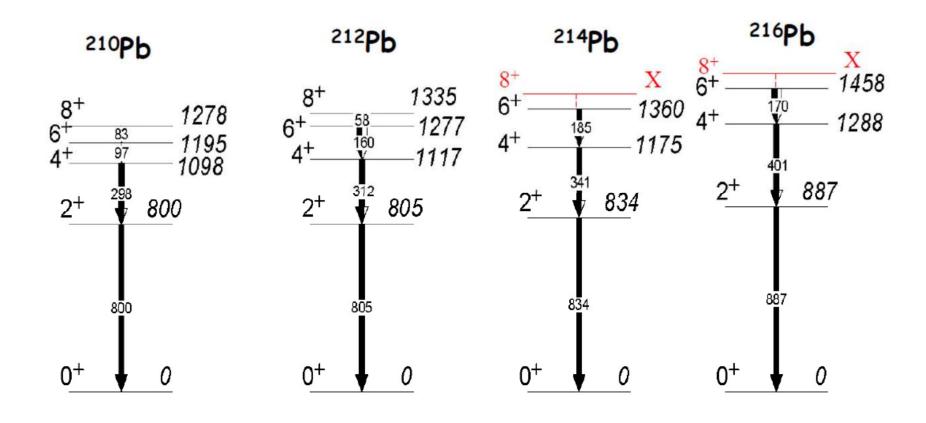
### <sup>212,214,216</sup>Pb: 8+ isomers:





### Energy levels well described in seniority scheme

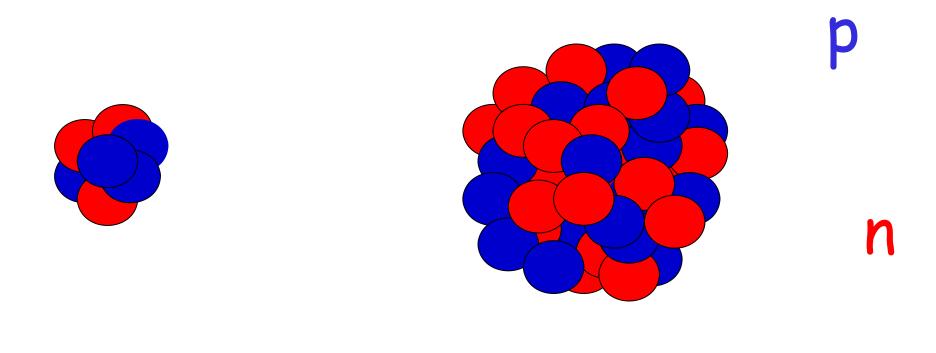
A. Gottardo, J.J. Valiente Dobon, G. Benzoni et al., PRL 109 (2012) 162502



# In-beam spectroscopy

```
Low-energy in-beam spectroscopy (up to 10 MeV per nucleon)
e.g.
Fusion evaporation.
Multi-nucleon transfer reactions
Coulomb excitation
Single nucleon transfer reactions
Intermediate-energy in-beam spectroscopy (50 -100 MeV per nucleon)
e.g.
Coulomb excitation
Spallation
Knockout
```

# E.g. fusion evaporation:



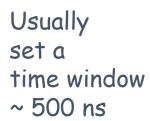
First particles are emitted..

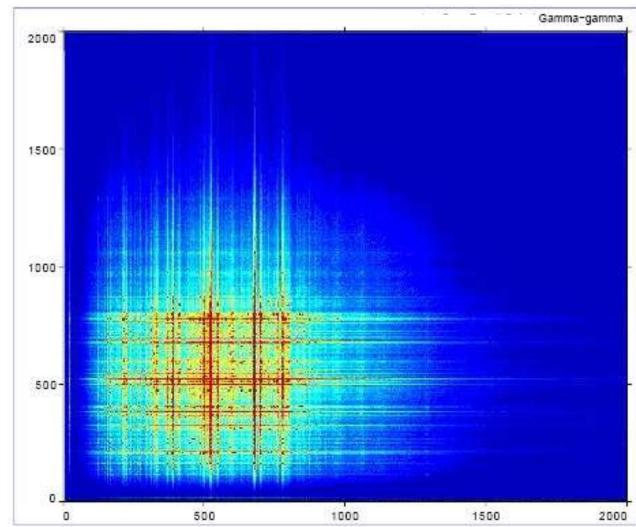
p

# Then gamma rays

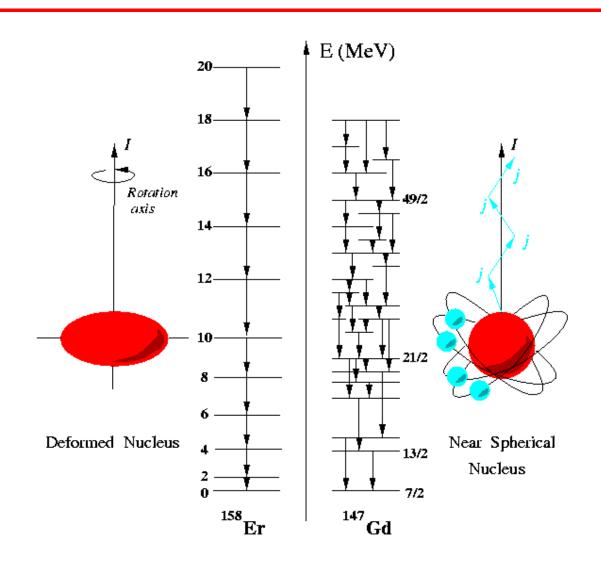
# **Level Schemes**

# Gamma-gamma coincidence matrices





# Gamma-ray patterns reveal nuclear structure



- Collective rotation (left) of a deformed nucleus leads to regular band structures
- Single-particle
  generation of spin
  (right) in a spherical
  nucleus leads to an
  irregular level structure

# Suggestions for tutorial discussion:

- 1. How can the 0<sup>+</sup> first excited state in <sup>40</sup>Ca decay to the ground state by 2 gamma rays if there are no intermediate levels?
- 2. Calculate E1, E2, M1, M2 transition rates for a single particle, 500 keV transition in an A=100 nucleus. What would the corresponding level lifetimes be?
- 3. What is the meaning of the phrase 'Energy levels well described in seniority scheme' when describing the Pb isomers? What particles are involved and in what orbits? (Pb, Z=82)
- 4. Estimate the angular momentum (in units hbar) brought into the compound nucleus  $^{156}$ Dy from the fusion of a  $^{48}$ Ca beam on a  $^{108}$ Pd target at a beam energy of  $E_{beam}$  = 206 MeV.

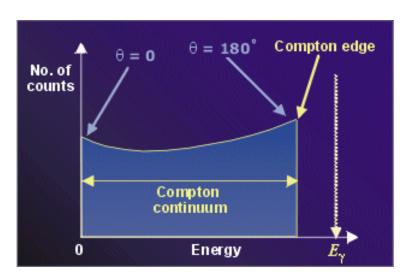
# Gamma-ray interactions in matter

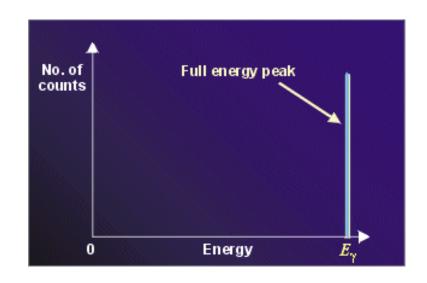
Gamma rays interact with matter via three main reaction mechanisms:

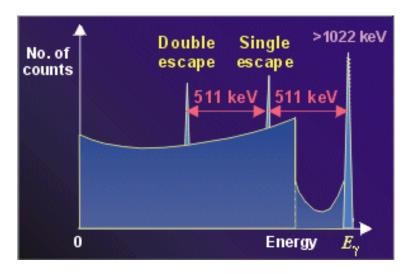
Photoelectric absorption

Compton scattering

Pair production





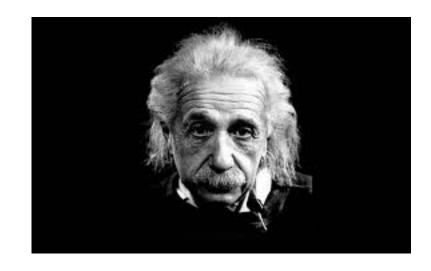


Pictures from University of Liverpool website

# Photoelectric absorption

Einstein won the Nobel Prize for Physics for the discovery of the photoelectric effect.

In this mechanism a y-ray interacts with a bound atomic electron.



The photon completely disappears and is replaced by an energetic photoelectron. The energy of the photoelectron can be written

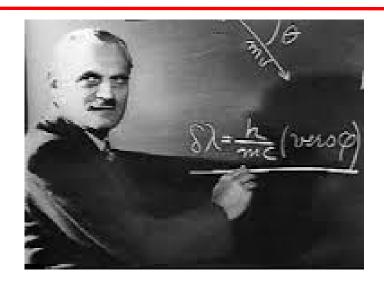
$$E_e = E_{\gamma} - E_{B}$$
.

The energy of the incident gamma-ray photon minus that of the binding energy of the electron ( $E_{\beta}$  = 12eV in germanium).

### Compton scattering

Compton won the Nobel Prize for Physics for the discovery of the Compton effect.

In this mechanism a  $\gamma$ -ray interacts with a loosely bound atomic electron.



The incoming  $\gamma$ -ray is scattered through an angle  $\theta$  with respect to its original direction.

The photon transfers a proportion of its energy to a recoil electron. The expression that relates the energy of the scattered photon to the energy of the incident photon is

$$E_{S} = \frac{E_{\gamma}}{1 + \frac{E_{\gamma}}{m_{\rho}c^{2}}(1 - \cos\theta)}$$

# Pair production

Nobody won the Nobel Prize for Physics for the discovery of the pair production effect (as far as I know).

If the energy of the  $\gamma$ -ray exceeds twice the rest mass energy of an electron (1.022 MeV), then the process of pair production is possible.





The incoming  $\gamma$ -ray disappears in the Coulomb field of the nucleus and is replaced by an electron-positron pair which has kinetic energy

 $E_{\gamma}$  - 1.022 MeV.

The positron is slowed down and eventually annihilates in the medium. Two annihilation photons are emitted back to back and these may or may not escape from the detector. Hence three peaks can be observed.

### Other interactions in a real detector

#### Thomson Scattering

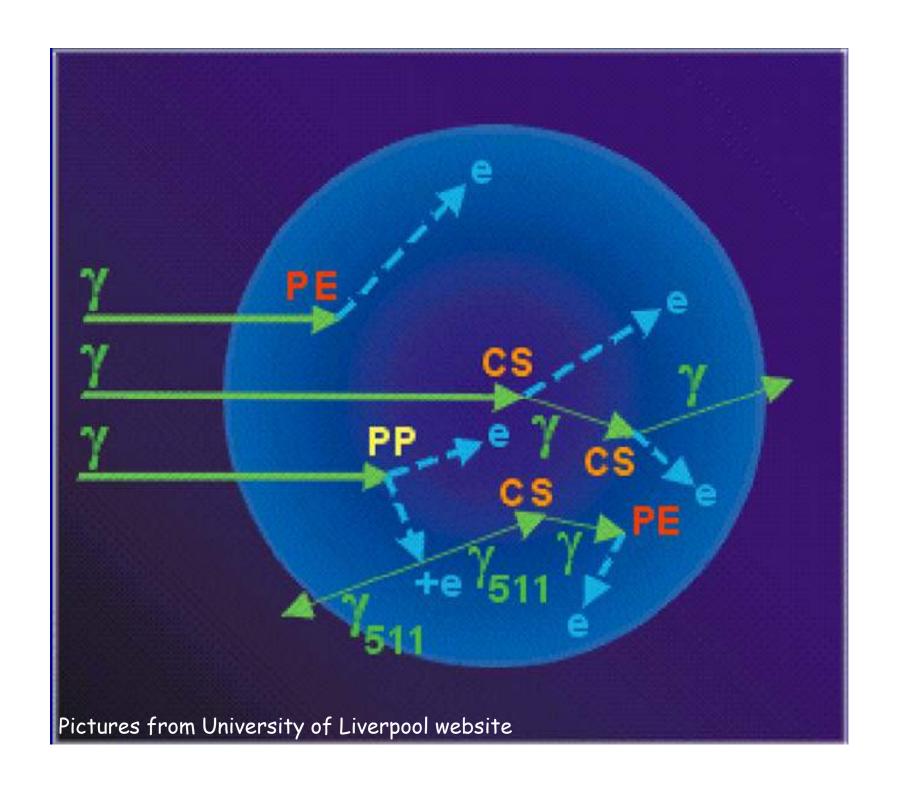
Low-energy coherent scattering off <u>free</u> electrons. Not important in the energy range concerned with most nuclear structure studies.

#### Nuclear Thomson Scattering

Low energy coherent scattering off nucleus. Small effect.

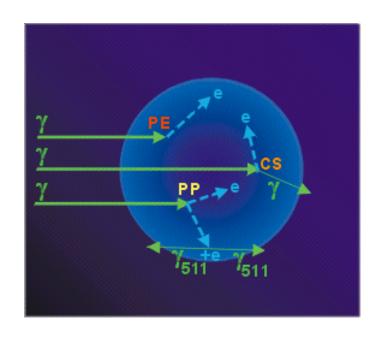
#### Dellbrück Scattering

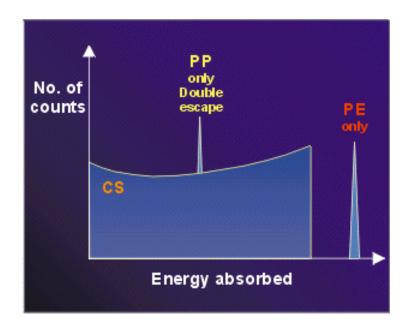
Scattering in the Coulomb field of the nucleus. Important at  $E_{\rm v}$  > 3 MeV.



#### Interactions in a small detector

A small detector is one in which only one interaction can take place. Only the photoelectric effect will produce full energy absorption. Compton scattering events will produce the Compton continuum. Pair production will give rise to the double escape peak due to both gamma-rays escaping.

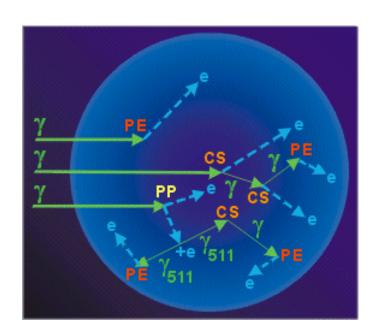


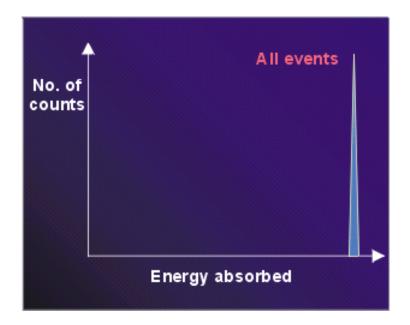


Pictures from University of Liverpool website

# Interactions in a large detector

A large detector is one in which there will be complete absorption of the gamma ray and a single gamma-ray peak, referred to as the full energy peak will be observed.





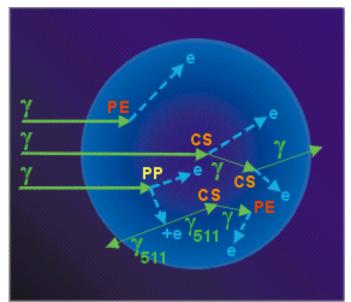
Pictures from University of Liverpool website

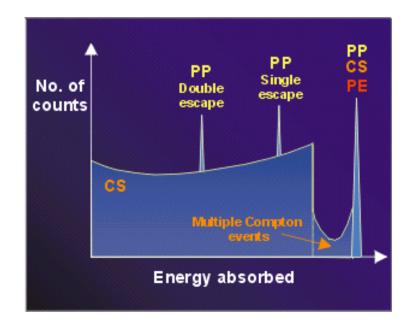
#### Interactions in a real detector

Within a real detector, the interaction outcome is not as simple to predict as e.g.

one Compton scattering could be followed by another before the gamma-ray photon escapes from the detector.

in the case of pair production, both, one or neither of the annihilation photons could escape from the detector. Hence all three peaks may be observed.



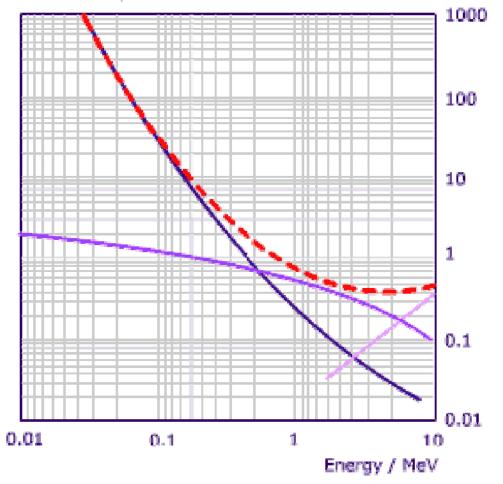


Pictures from University of Liverpool website

# Energy dependence of the interactions



Linear attenuation / cm<sup>-1</sup>



Typically we are interested in transitions of energy  $60 \text{ keV} < E_{\gamma} < 10 \text{ MeV}.$ 

### Detector types

Two main types of material are used:

Solid state semiconductor detectors e.g. Ge, CZT

Electron-hole pairs are collected as charge

knock-on effect => an avalanche arrives at the electrode

lots of electrons => good energy resolution

cooled to liquid  $N_2$  temperature (77K) to reduce noise

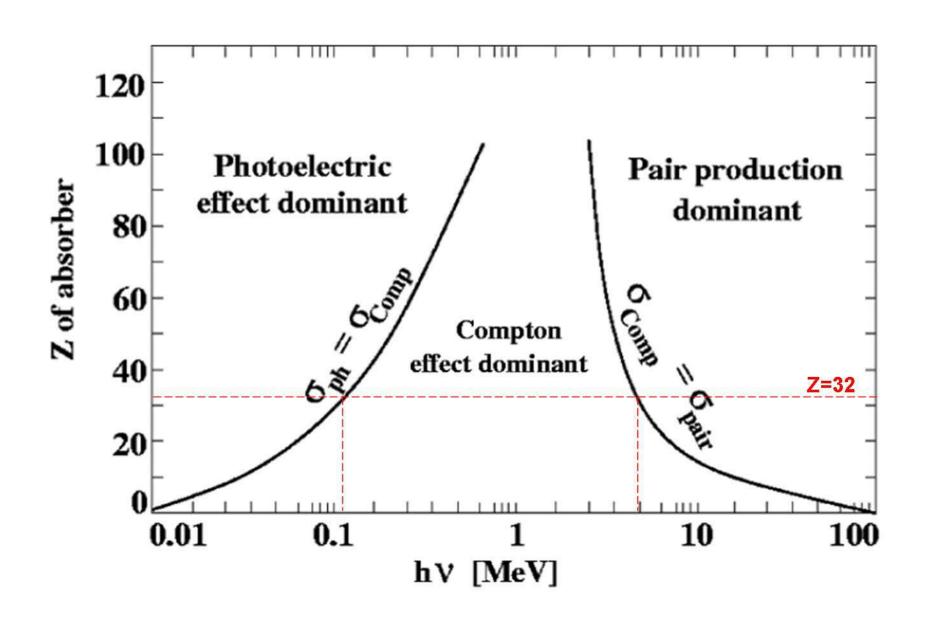
Advantage: good energy resolution (~0.15 % (FWHM) at 1.3 MeV)

Disadvantages: relatively low efficiency

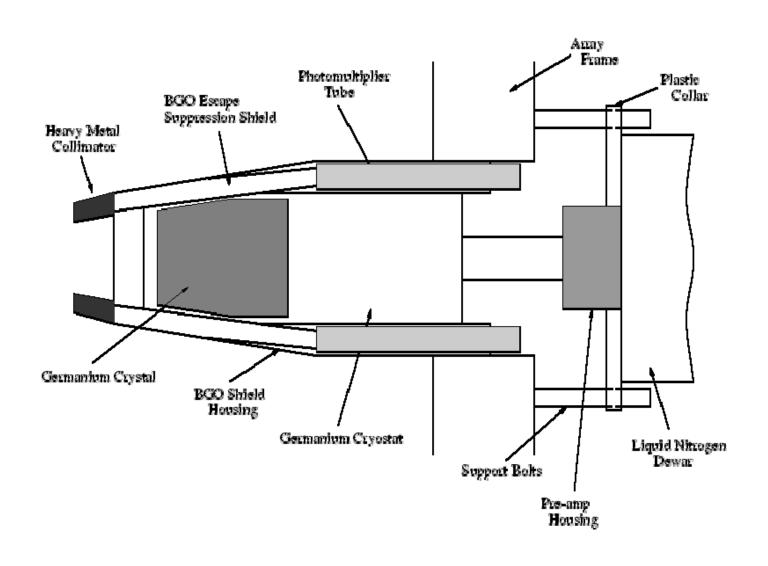
cryogenic operation

limit to the size of crystal/detector

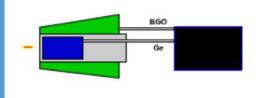
# Z dependence of interaction probablities

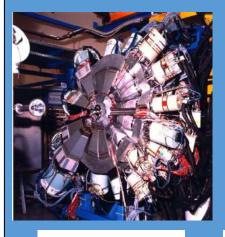


# Escape suppressed HP Ge detector



Large Gamma Arrays based on Compton Suppressed Spectrometers





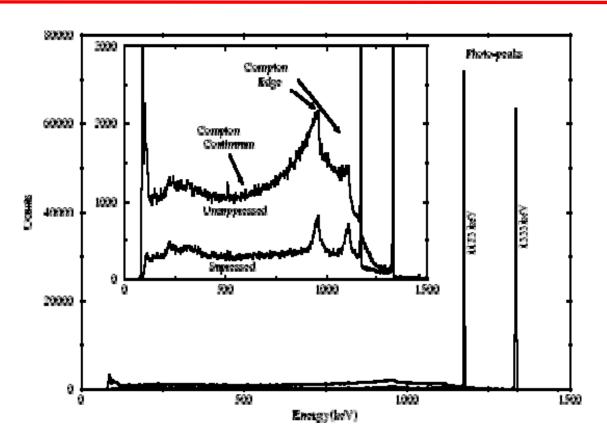


**EUROBALL** 

**GAMMASPHERE** 

$$\varepsilon \sim 10 - 5 \%$$
(  $M_{\gamma}=1 - M_{\gamma}=30$ )

### Compton suppression increases peak to total



Bare detector: P/T ~20% at ~ 1 MeV i.e only 20% of single events have the full energy measured for  $\gamma-\gamma$  only 4% of events are full energy,  $\gamma-\gamma-\gamma-\gamma$  0.16%

Compton suppression: P/T ~ 60%

### Detector types

Two main types of material are used:

Scintillation detectors e.g. NaI, BGO, LaBr<sub>3</sub>(Ce)

Recoiling electrons excite atoms, which then de-excite by emitting visible light.

Light is collected in photomultiplier tubes (PMT) where it generates a pulse proportional to the light collected.

Advantage: good timing resolution

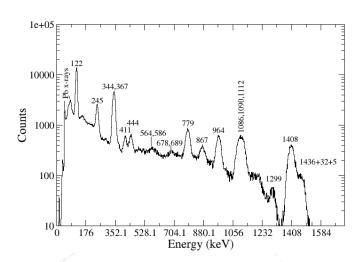
can be made relatively large e.g. NaI detectors 14"  $\phi$  x 10" no need for cryogenics

Disadvantages: poor energy resolution

#### Scintillation detectors

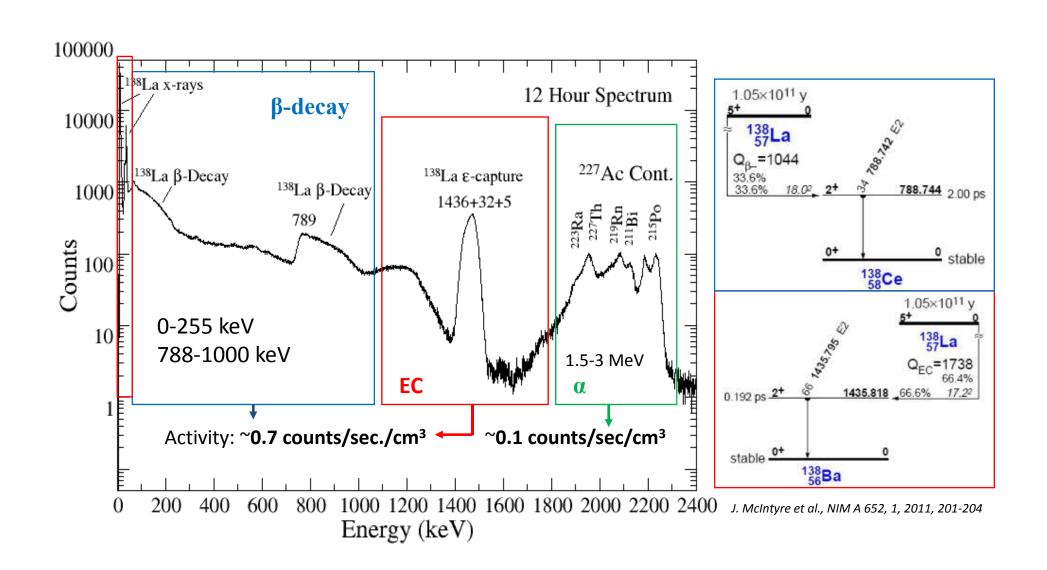
#### $LaBr_3(Ce)$

- LaBr $_3$ (Ce) timing properties:
  - ~ 25 ns decay time
  - Timing Resolution FWHM of 130-150 ps with 60Co for a Ø1"x1" crystal.
- High energy resolution, 3 % FWHM at 662 keV.
- Peak Emission wavelength in Blue/UV part of EM spectrum (380 nm), compatible with PMTs.





# Detector Characterisation



# Timing resolution of cylindrical crystals



ø1"x1"

FWHM 200 ps

#### FWHM 150 ps

N. Mărginean et al. Eur. Phys. J A 46, 329-336, 2010.

I. Deloncle et al. J. Phys.: Conf. Series 205, 012044, 2010.

M. Moszynski et al. Nucl. Instr. Methods A 567, 2007.

L.M. Fraile et al. ISOLDE Workshop, Fast timing results at ISOLDE, http://indico.cern.ch/getFile.py/access?contribId= 36&sessionId=8&resId=0&materialId=slides&confId=67060, November, 2009.



ø1.5"x1.5"

360 ps

180 ps



ø2"x2"

450 ps at 511 keV 300 ps at 1332 keV

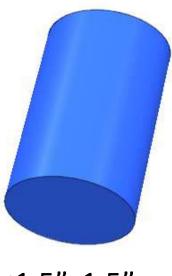
# Timing resolution of cylindrical crystals



ø1"x1"

FWHM 200 ps

FWHM 150 ps



ø1.5"x1.5"

360 ps

180 ps



ø2"x2"

450 ps at 511 keV

300 ps at 1332 keV

Trade off between resolution and efficiency

# Timing Precision

Trade off between good timing resolution (small detectors)

lots of statistics (large detectors)

Timing precision defined as:

TP= 
$$\frac{T(FWHM)}{\sqrt{N}}$$

# Detector requirements for in-beam spectroscopy

#### Gamma rays emitted by moving reaction product:

Good energy resolution (as lots of gamma rays emitted)

High granularity (to uniquely define the  $\gamma$ -ray angle and because lots of gamma rays are emitted – high multiplicity)

High photopeak detection efficiency (to see the weakest channels)

Good peak to total ratio (so that coincidence gates can be clean)

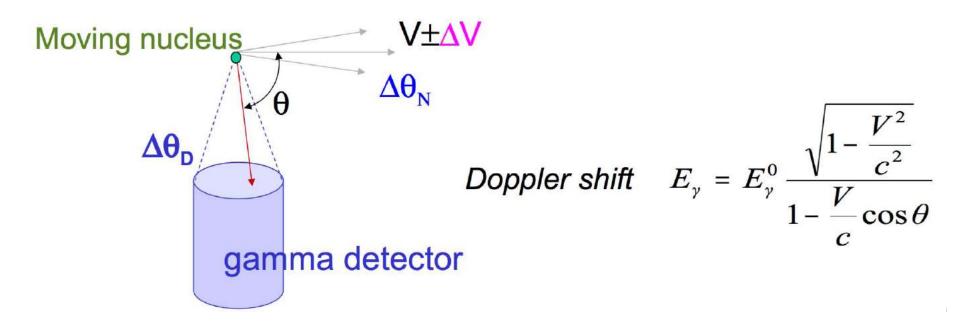
# Energy resolution

The major factors affecting the final energy resolution (FWHM) at a particular energy are as follows:

$$\Delta E_{\gamma}^{\text{final}} = \left(\Delta E_{\text{Int}}^2 + \Delta \theta_{\text{D}}^2 + \Delta \theta_{\text{N}}^2 + \Delta V^2\right)^{1/2}$$

- $\Delta E_{\rm Int}$  The intrinsic resolution of the detector system. This includes contributions from the detector itself and the electronic components used to process the signal.
- $\Delta\theta_{\text{D}}$  The Doppler broadening arising from the opening angle of the detectors.
- $\Delta\theta_{N}$  The Doppler broadening arising from the angular spread of recoils in the target.
- $\Delta V$  The Doppler broadening arising from the velocity (energy) variation of the recoils across the target.

# Doppler broadening



Broadening of detected  $\gamma$ -ray energy arises from:

- Spread in recoil velocity △V
- •Distribution in the direction of recoil  $\Delta \theta_{N}$
- •Detector opening angle  $\Delta \Theta_{D}$

# Minimising Doppler broadening

There are two ways in which gamma-ray spectroscopists can mitigate the effects of Doppler broadening.

- •Reduce the detector opening angle (lower  $\Delta\theta_D$ )
  - Detector granularity
- •Minimise the target thickness (lower spread in recoil velocity  $\Delta V$ )

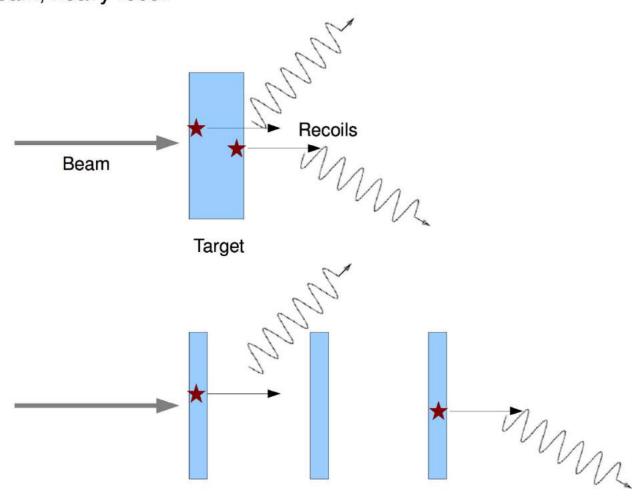
Experimentalists often choose to use two or three stacked targets rather than a single thick target e.g.  $2 \times 0.5 \text{ mg/cm}^2$  rather than a single  $1 \text{ mg/cm}^2$ .

This works for normal kinematics where the spread in velocities  $\Delta V$  arises from the slowing of the recoil and not the beam.

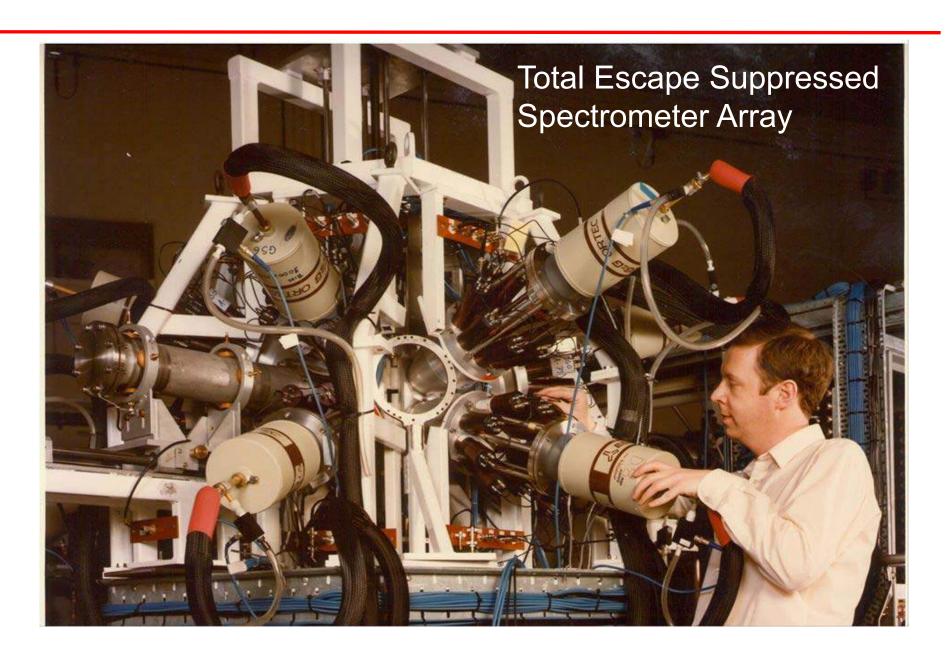
# Minimising Doppler broadening

#### **Target thickness**

Light beam, heavy recoil

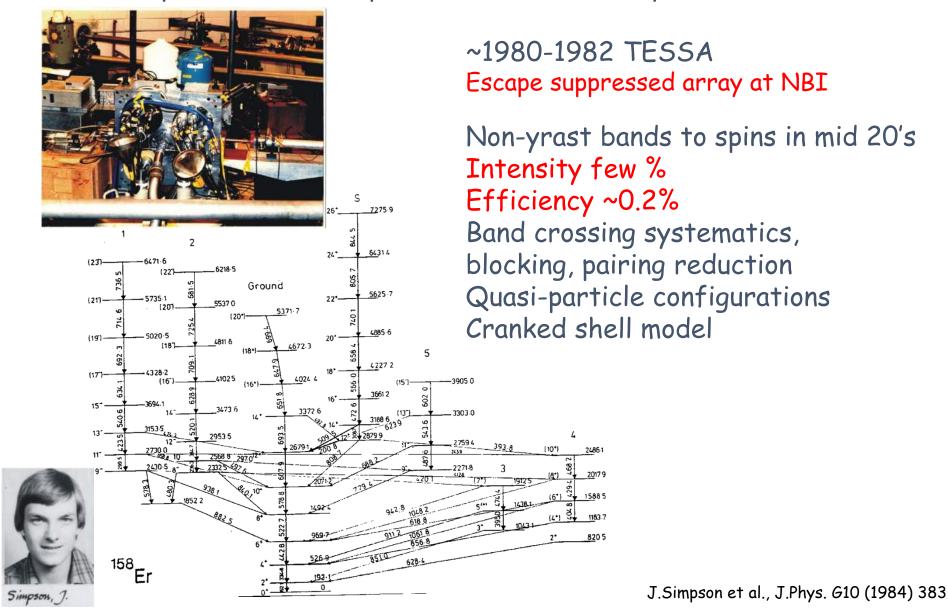


# Successful Compton suppression arrays:

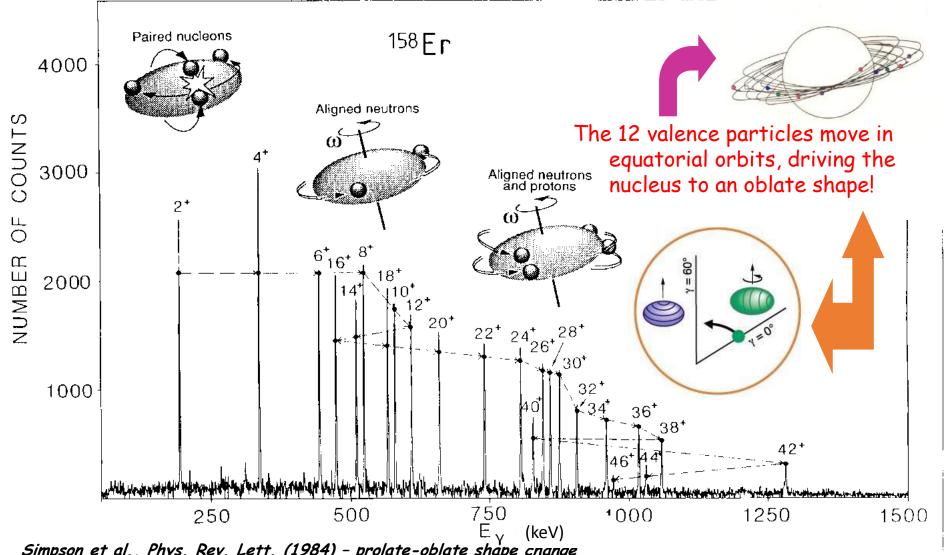


#### Spectroscopy of 158Er

~1980 yrast states to spin ~30, naked Ge arrays



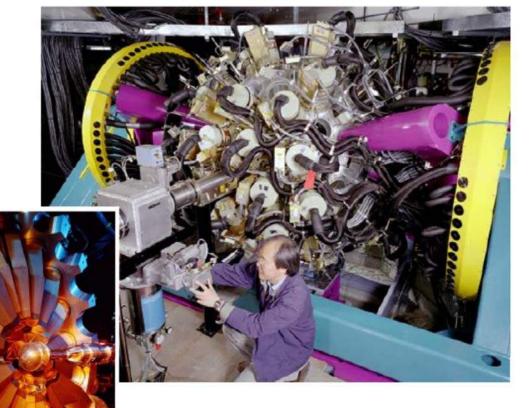
#### <sup>158</sup>Er expt. Daresbury -Sharpey-Schafer, Riley, Simpson et al. mid 1980's



Simpson et al., Phys. Rev. Lett. (1984) - prolate-oblate shape cnange P.O. Tjom et al., PRL 55 (1985) 2405 - lifetime measurements T. Bengtsson and I. Ragnarsson, Physica Scripta T5 (1983) 165 J. Dudek, W. Nazarewicz Phys. Rev C32 (1985) 298 Ragnarsson, Xing, Bengtsson and Riley, Phys. Scripta 34 (1986) 651

# Successful Compton suppression arrays:

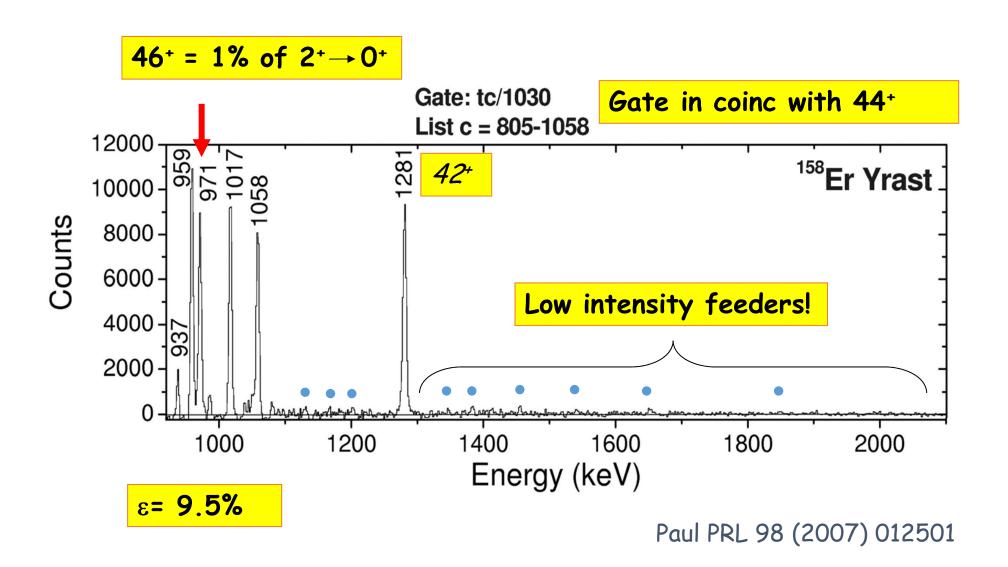
#### Gammasphere



110 Compton-suppressed Ge detectors, each with 70% efficiency. Total efficiency = 9% at 1.3 MeV.

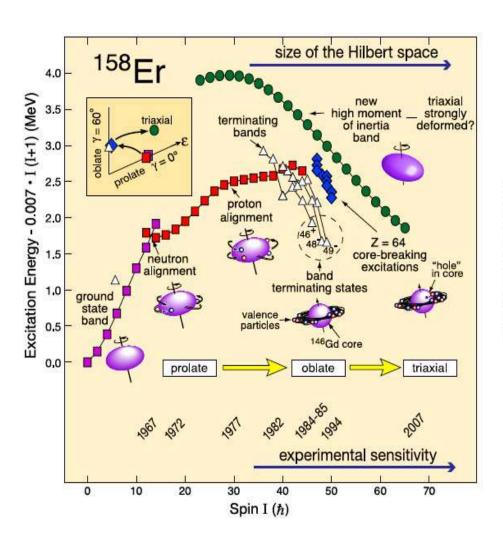
# What about <sup>158</sup>Er above 46<sup>+</sup>?

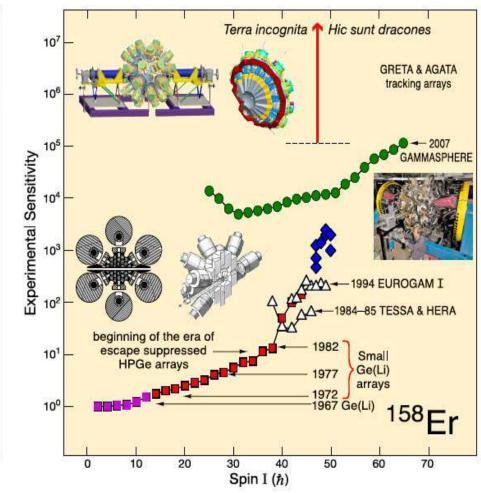
No wonder we could not see it before!



# Evolution of Gamma-Ray Spectroscopy

### New Detector Systems ← New Physics

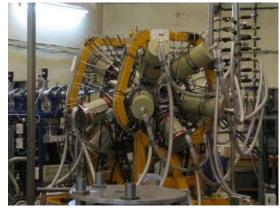




# Mixed arrays

More recently,  $LaBr_3(Ce)$  detectors have been added to arrays of Compton suppressed detectors because of their good timing properties.

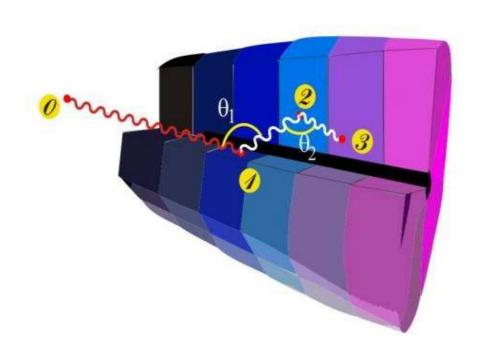
#### E.G. ROSPHERE at Bucharest





- 15 HPGe detectors (A/C):
  - 10 x HPGe detectors @ 37°
  - 1 x HPGe detector @ 64°
  - 4 x HPGe detectors @ 90°
- 11 LaBr<sub>3</sub>(Ce):
  - ø2"x2" @ 90 and 64° (three)
     (Cylindrical)
  - ø1.5"×1.5" @ 90 (six) (Cylindrical)
  - ø1"x1.5" @ 64° (two) (Conical)

# Next generation: tracking

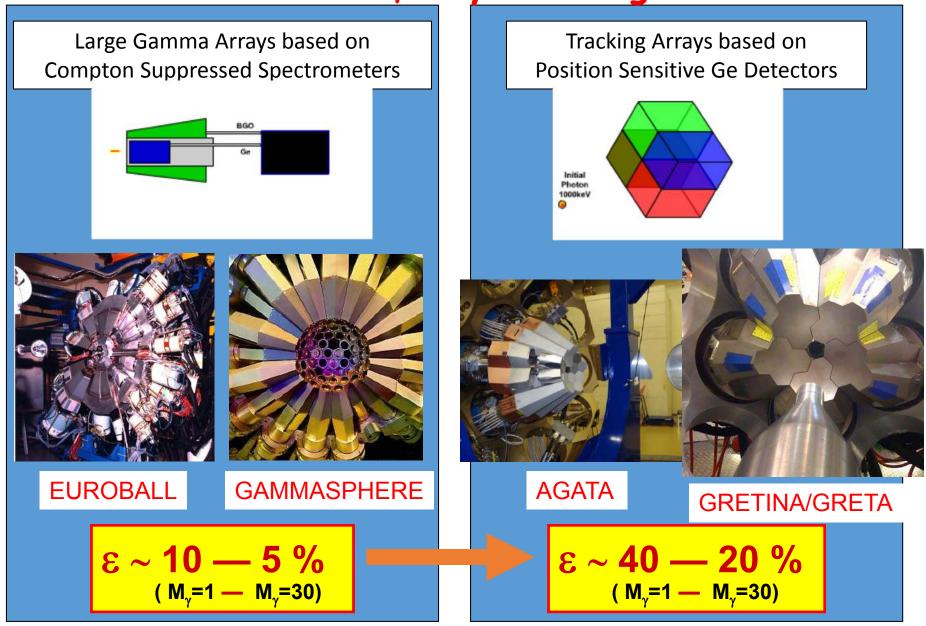


Detectors are segmented

longitudinally and radially.

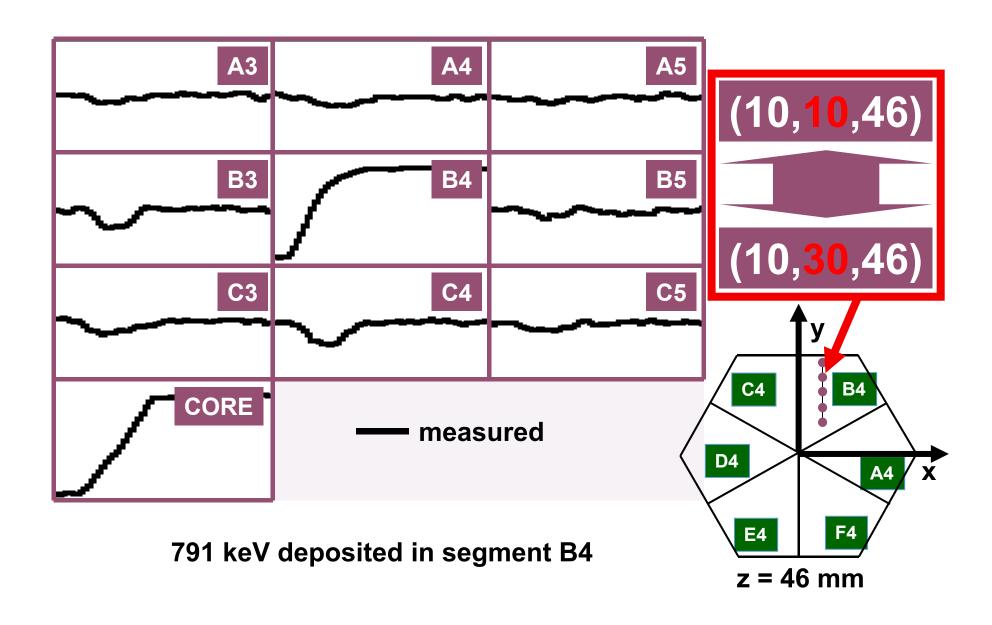
Pinpoint the position of the first interaction to get the angle of the incident gamma ray (very important if have a high v/c).

Idea of  $\gamma$ -ray tracking

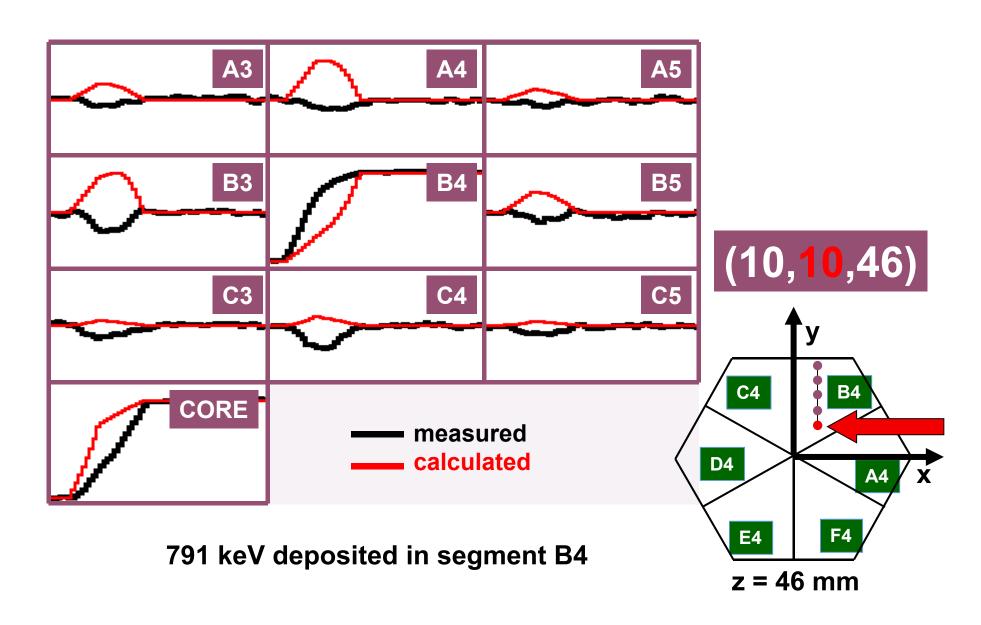


Huge increase in sensitivity

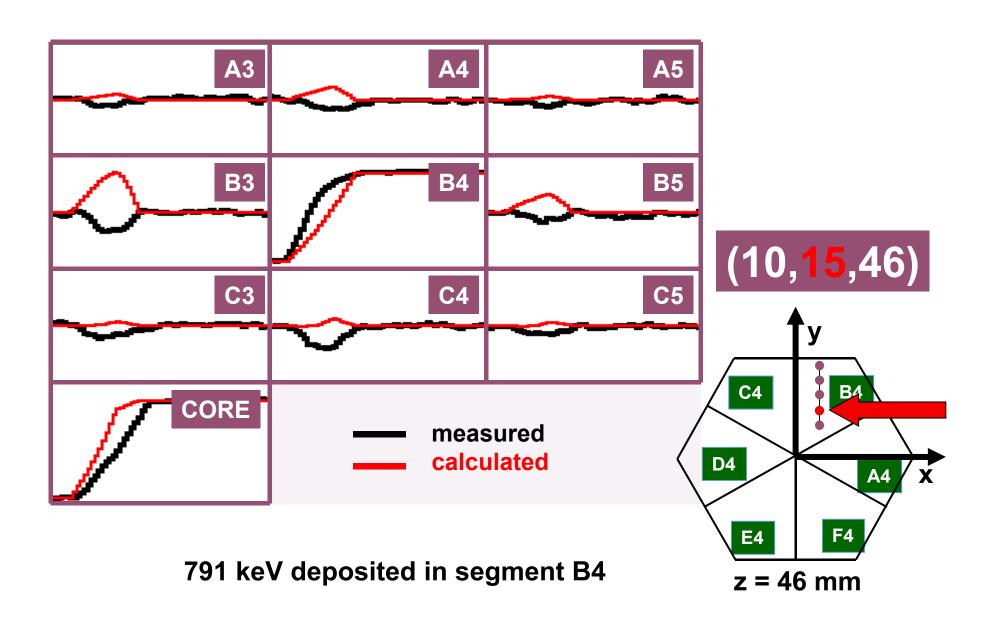
# Pulse Shape Analysis concept

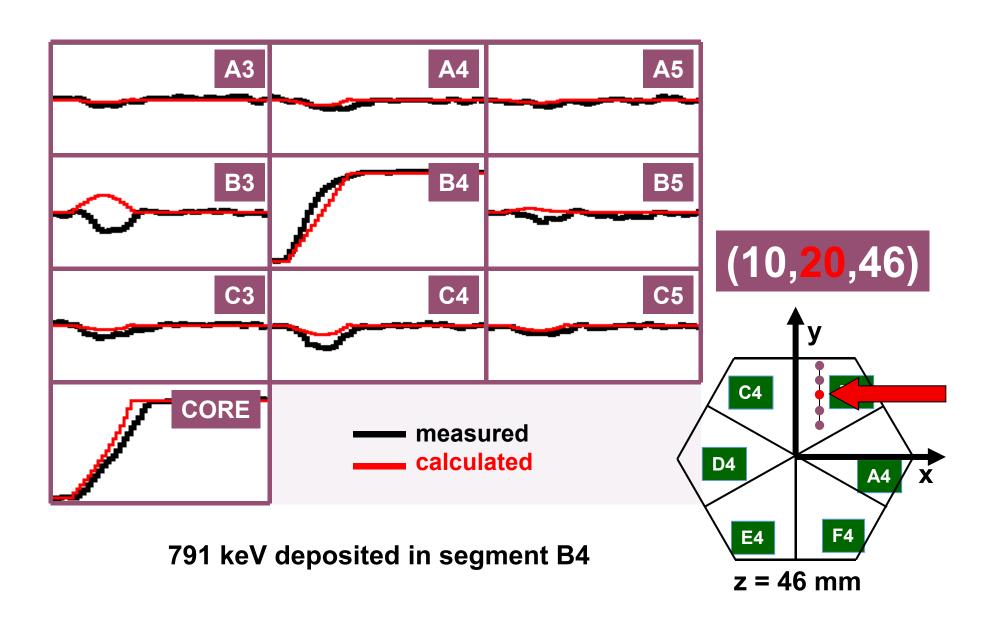


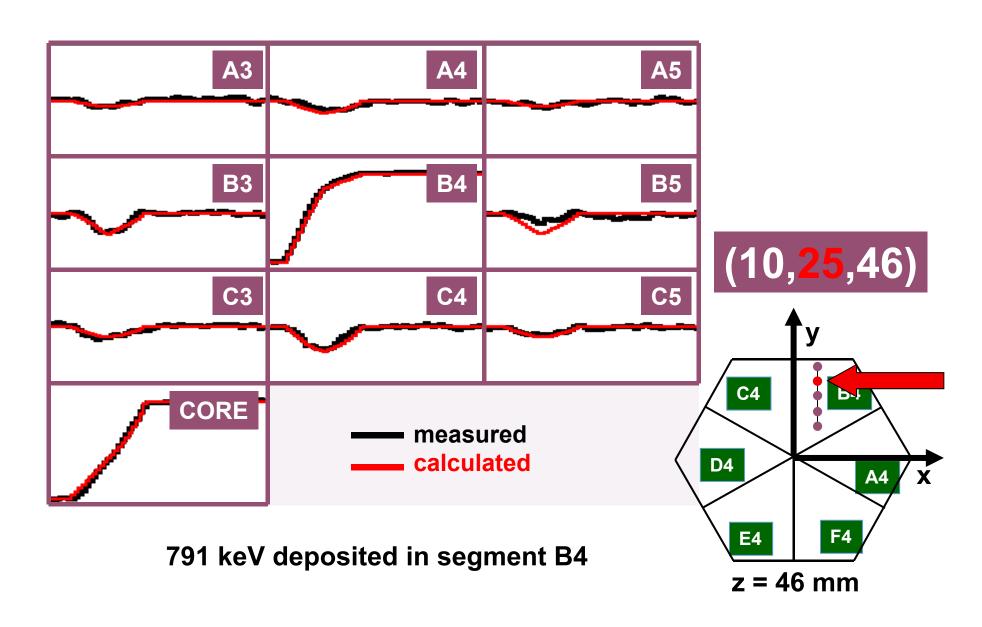
# Pulse Shape Analysis concept

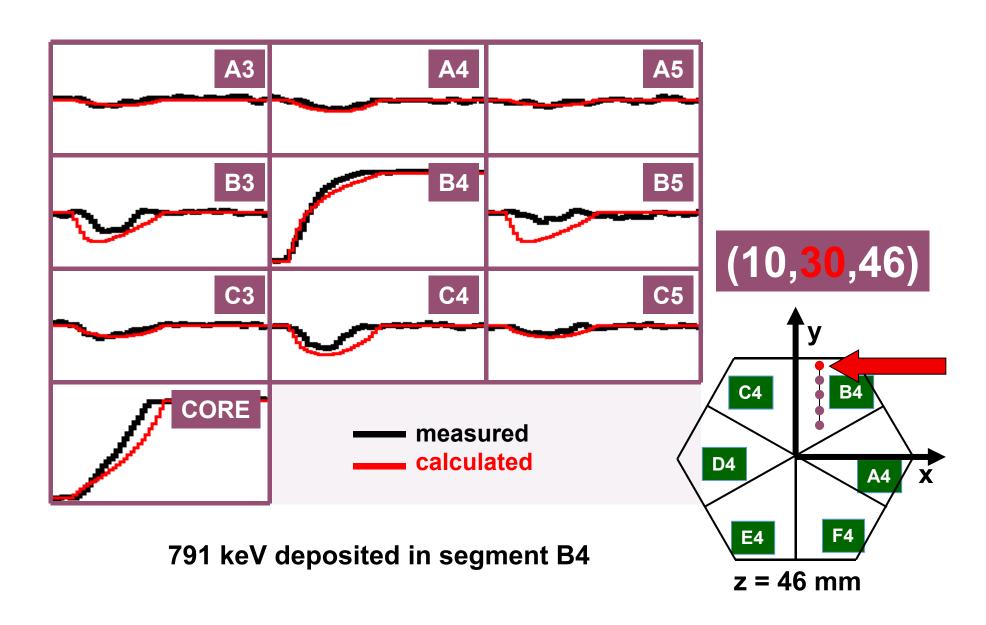


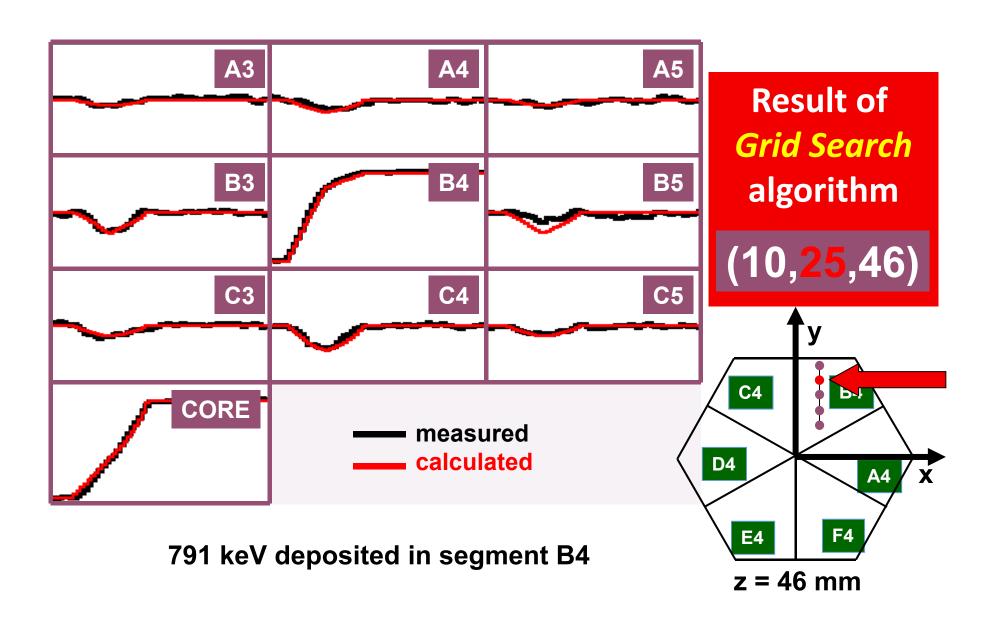
# Pulse Shape Analysis concept







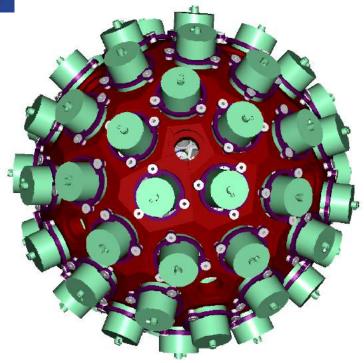




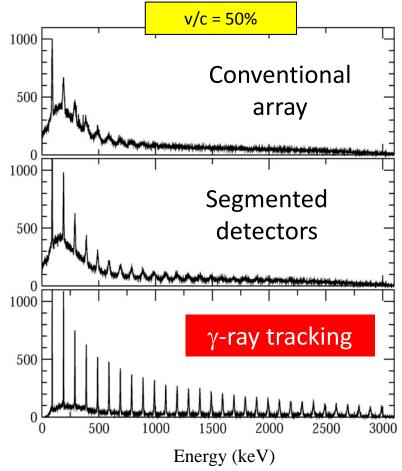


## AGATA (Advanced GAmma Tracking Array)



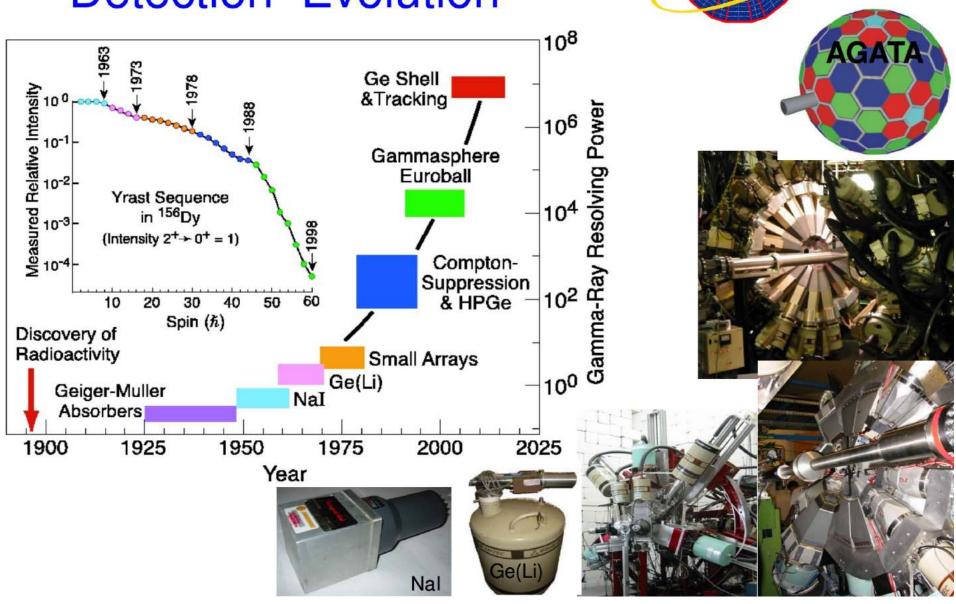


The innovative use of detectors (pulse shape analysis,  $\gamma$ -ray tracking, digital DAQ) will result in high efficiency (~40%) and excellent energy resolution, making AGATA the ideal instrument for spectroscopic studies of weak channels.



The effective energy resolution is maintained also at "extreme" v/c values

# Gamma-Ray Detection Evolution



## Detector requirements for out-of-beam spectroscopy

#### Gamma rays emitted by a stopped source:

Doppler broadening is not an issue..detectors can be very close but if you do this you lose angle definition which is required for angular measurements.

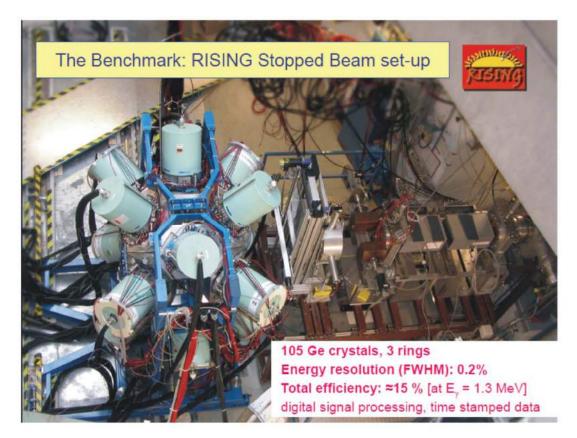
Few gamma rays means you can have fewer big detectors but see note above re detector angle

High photopeak detection efficiency (to see the weakest channels)

Good peak to total ratio (so that coincidence gates can be clean)

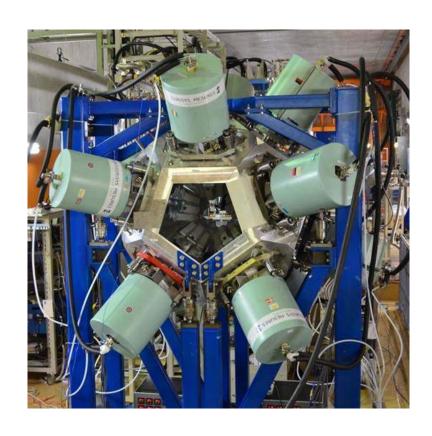
## Detector requirements for out-of-beam spectroscopy

In recent years, experiments have still tended to be performed with arrays e.g. the use of the RISING array of Euroball clusters for experiments at the FRS at GSI.



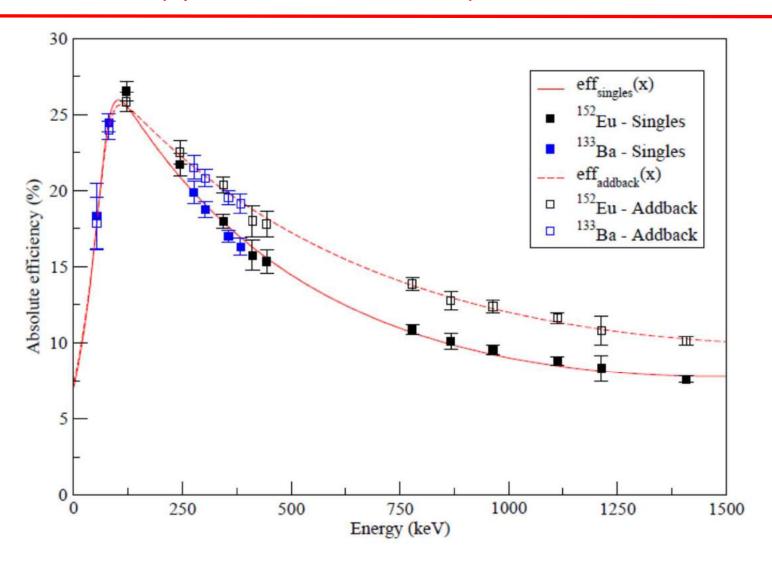
## Detector requirements for out-of-beam spectroscopy

Now also using mixed arrays e.g. the use of the EURICA array of 12 Euroball clusters with 18 LaBr $_3$ (Ce) detectors at BigRIPS at RIKEN.





## Typical efficiency curve



Efficiency of the 12 EURICA Cluster detectors for singles (solid symbols) and add back (open symbols)

## Suggestions for tutorial discussion:

- 1. Explain the origin of the part of the spectrum labelled "multiple Compton events" in the spectrum for the 'real detector'
- 2. Why are Compton suppression shields made from scintillator material?
- 3. Why is the timing resolution worse for bigger scintillator detectors?
- 4. Why does the efficiency curve have this shape?

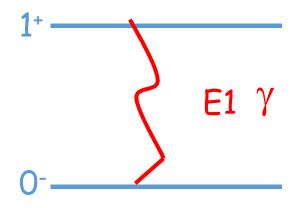
### Spins and parities

Two distinct types of measurements:

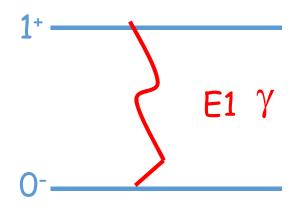
Angular correlation: can be done with a non-aligned source but need  $\gamma-\gamma$  coincidence information.

Angular distribution: need an aligned source but can be done with singles data.

...note that these cannot measure parity but you can usually infer something about the transition.



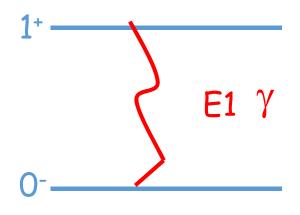
Imagine the situation of an E1 decay between two states, the initial one has a  $J^{\pi}$  value of 1<sup>+</sup> and the final one a  $J^{\pi}$  of 0<sup>-</sup>.



The  $J^{\pi} = 1^+$  state has 3 substates with m values of  $\pm 1$  and 0.

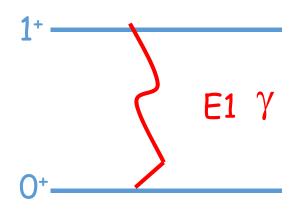
The  $J^{\pi} = 0^{-}$  state has only 1 substate, with m=0.

When the substates of the  $J^{\pi}$  = 1<sup>+</sup> state decay, the  $\gamma$  rays emitted have different angular patterns.



$$J^{\pi} = 1^+$$
, m=0 decays to  $J^{\pi} = 0^-$ , m=0 with a  $\sin^2\theta$  distribution.

```
J^{\pi} = 1^+, m=±1 decays to J^{\pi} = 0^-, m=0 with a \frac{1}{2}(1+\cos^2\theta) distribution.
```

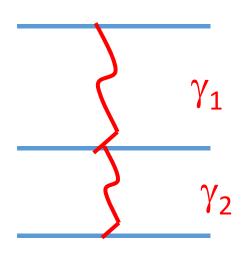


$$J^{\pi} = 1^+$$
, m=0 decays to  $J^{\pi} = 0^+$ , m=0 with a  $\sin^2\theta$  distribution.

$$J^{\pi} = 1^+$$
, m=±1 decays to  $J^{\pi} = 0^+$ , m=0 with a  $\frac{1}{2}(1+\cos^2\theta)$  distribution.

So the total distribution is 
$$\frac{1}{2}(1+\cos^2\theta) + \sin^2\theta + \frac{1}{2}(1+\cos^2\theta)$$
  
=  $1+\cos^2\theta + \sin^2\theta$   
= 2 ...flat, no angular dependence

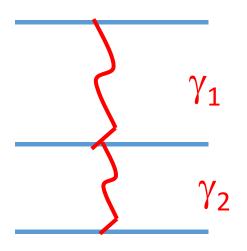
### Angular correlation - non-oriented source



Let's imagine we have two  $\gamma$ rays which follow
immediately after each
other in the level scheme.

If we measure  $\gamma_1$  or  $\gamma_2$  in singles then the distribution will be isotropic (same intensity at all angles)... there is no preferred direction of emission.

#### Angular correlation - non-oriented source



Now imagine that we measure  $\gamma_1$  or  $\gamma_2$  in coincidence. We say that measuring  $\gamma_1$  causes the intermediate state to be aligned. We define the z direction as the direction of  $\gamma_1$ 

The angular distribution of the emission of  $\gamma_2$  then depends on the spin/parities of the states involved and on the multipolarity of the transition.

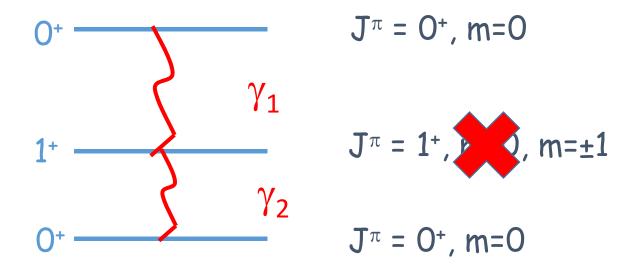
## A simple example:

$$O^{+}$$
  $J^{\pi} = O^{+}, m=0$ 

$$J^{\pi} = 1^{+}, m=0, m=\pm 1$$

$$J^{\pi} = O^{+}, m=0$$

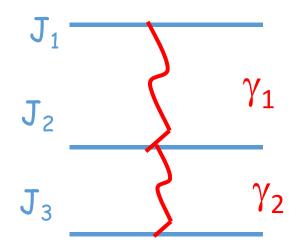
## A simple example:



Hence for  $\gamma_2$  we only see the m =  $\pm$  1 to m=0 part of the distribution i.e we see that the intensity measured as a function of angle (w.r.t  $\gamma_1$ ) follows a  $1+\cos^2\theta$  distribution.

#### General formula

In general,
The gamma-ray intensity
varies as:



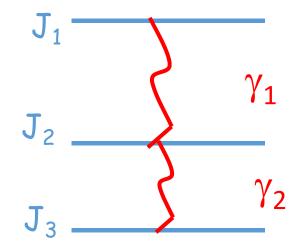
$$W(\theta) = \sum_{k_{even}} A_k (\gamma_1) A_k (\gamma_2) Q_k (\gamma_1) Q_k (\gamma_2) P_k (\cos \theta)$$

where

 $\theta$  is the relative angle between the two  $\gamma$ -rays  $Q_k$  accounts for the fact that we do not have point detectors

 $A_k$  depends on the details of the transition and the spins of the levels

#### General formula



$$A_k(\gamma_1) = \frac{F_k(J_2J_1ll) - 2\delta F_k(J_2J_1ll + 1) + \delta^2 F_k(J_2J_1l + 1l + 1)}{1 + \delta^2}$$

$$A_k(\gamma_2) = \frac{F_k(J_2J_3LL) + 2\delta F_k(J_2J_3LL + 1) + \delta^2 F_k(J_2J_3L + 1L + 1)}{1 + \delta^2}$$

The  $F_k$  coefficients contain angular momentum coupling information .....3j, 6j symbols.

## Legendre polynomials

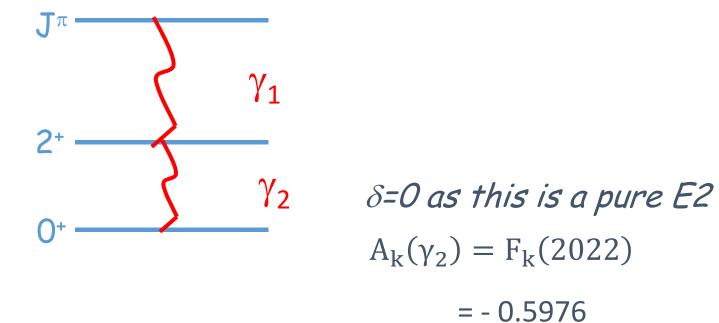
$$P_2(\cos\theta) = \frac{1}{2} (3\cos^2\theta - 1)$$

$$P_4(\cos\theta) = \frac{1}{8} (35\cos^4\theta - 30\cos^2\theta + 3)$$

$$P_6(\cos\theta) = \frac{1}{16} (231\cos^6\theta - 315\cos^4\theta + 105\cos^2\theta - 5)$$

Note the dependence on cos<sup>2</sup>.

## A specific case:



### A specific case:

<sup>195</sup>Pt( $n,\gamma$ )<sup>196</sup>Pt reaction

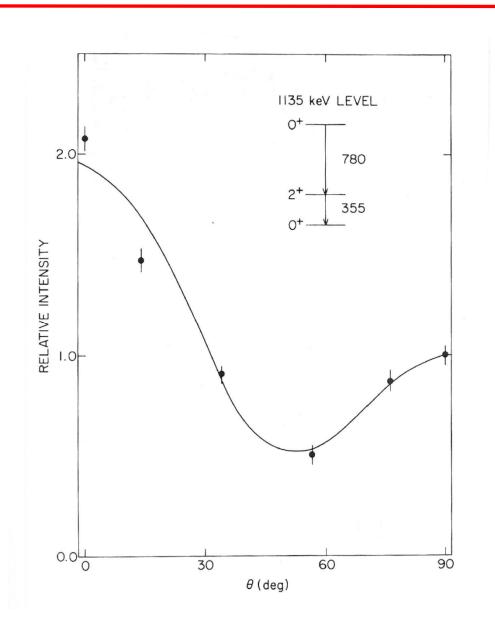
slow-neutron capture so definitely no alignment brought into the system

v simple system, 2 detectors 1 moving 1 fixed

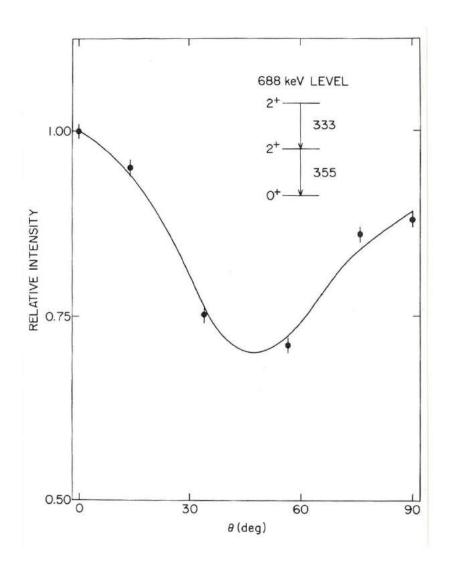
circa. 1983

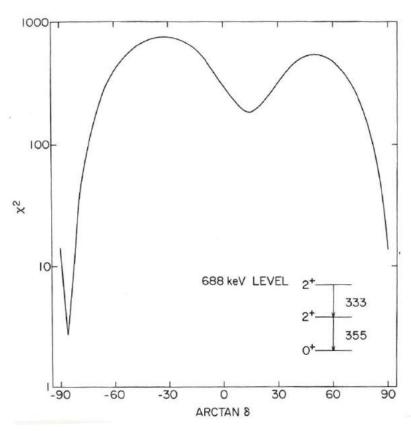


## A specific case: $^{195}Pt(n,\gamma)^{196}Pt$



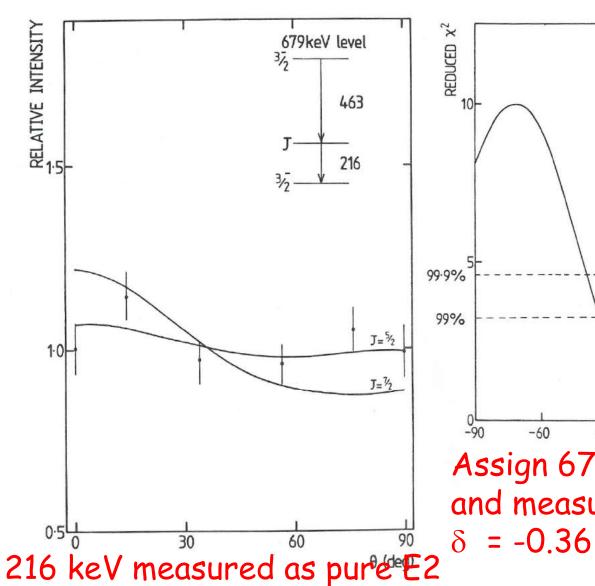
## A specific example - extract mixing ratios

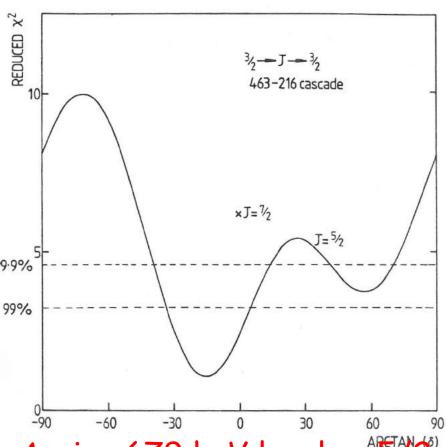




arctan 
$$\delta \sim -86$$
  
 $\delta = -14.3$  (40)

## A specific example - assign spins 188Os(n,y)189Os





Assign 679 keV level as  $572^{\circ}$  and measure arctan  $\delta \sim -20$   $\delta = -0.36$  12% quadrupole 88% dipole

## Angular correlations with arrays

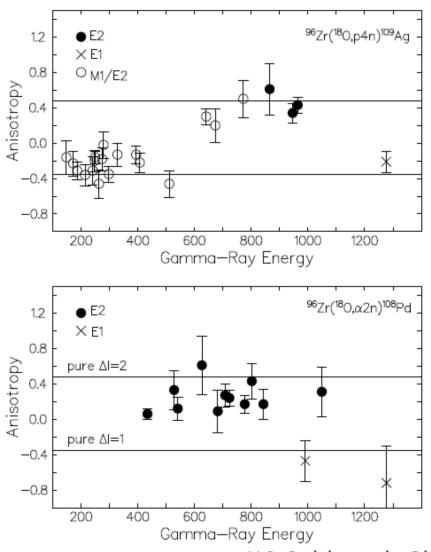
Many arrays are designed symmetrically so the range of possible angles is reduced.

In these cases it is most common to measure a 'DCO' ratio. e.g. in the simplest case, if you have an array with detectors at 35° and 90°: Gate on 90° detector, measure coincident intensities in:

- other 90° detectors
- 35° detectors

Take the ratio and compare with calculation...can usually separate quadrupoles from dipoles but cannot measure mixing ratios.

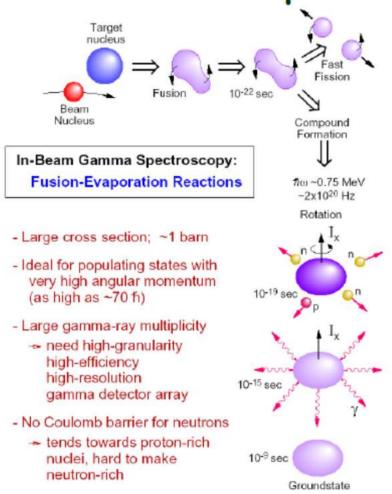
## Angular correlations with arrays

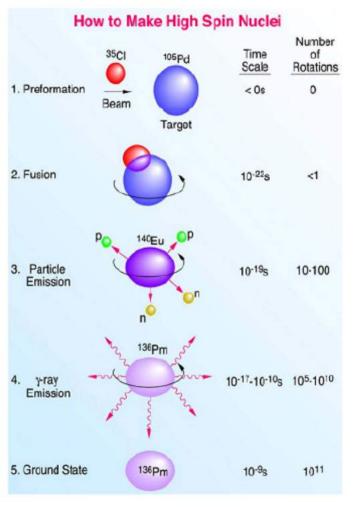


K.R.Pohl et al., Phys Rev C53 (1996) 2682

## Angular distribution

## Fusion Evaporation Reactions





### Angular distribution

In heavy-ion fusion-evaporation reactions, the compound nuclei have their spin aligned in a plane perpendicular to the beam axis:

$$\ell = r \times p$$

Depending on the number and type of particles 'boiled off' before a  $\gamma$  ray is emitted, transitions are emitted from oriented nuclei and therefore their intensity shows an angular dependence.

## Angular distribution

$$W(\theta) = A_0 + A_2 B_2 P_2(\cos \theta) + A_4 B_4 P_4(\cos \theta) + A_6 B_6 P_6(\cos \theta) + \cdots$$

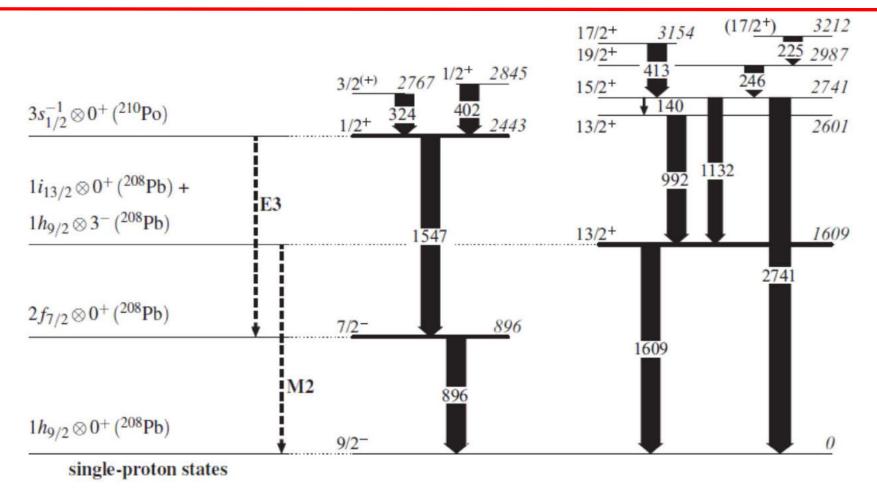
$$= A_0(1 + \frac{A_2}{A_0}B_2P_2(\cos\theta) + \frac{A_4}{A_0}B_4P_4(\cos\theta) + \frac{A_6}{A_0}B_6P_6(\cos\theta) + \cdots)$$

$$= A_0(1 + a_2B_2P_2(\cos\theta) + a_4B_4P_4(\cos\theta) + a_6B_6P_6(\cos\theta) + \cdots)$$

Where  $A_k$  and  $P_k$  are as before and  $B_k$  contains information about the alignment of the state

(in principle there should be a  $Q_k$  in here too but let's forget for now)

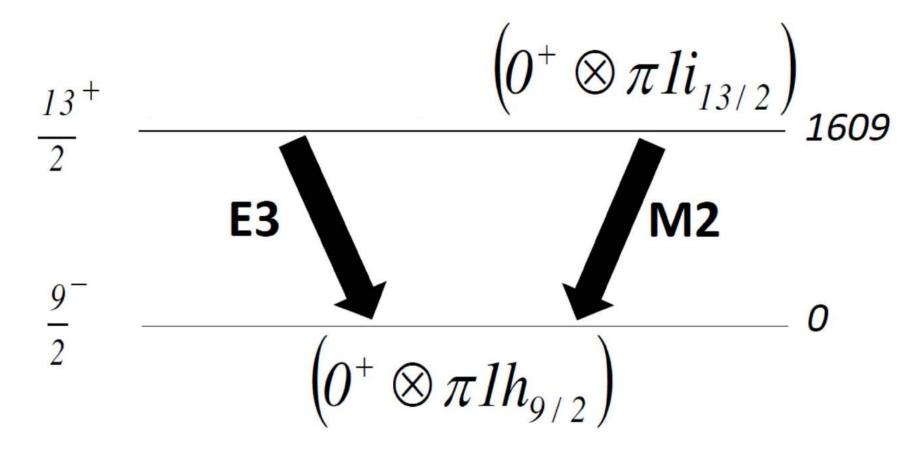
## Angular distribution: worked example 209Bi



Measure mixing ratio of 1609 keV transition

## Angular distribution: worked example 209Bi

#### What's the physics?



## Angular distribution: worked example 209Bi

<sup>209</sup>Bi was populated in the <sup>208</sup>Pb( $^{7}$ Li,2n $\alpha\gamma$ )<sup>209</sup>Bi

reaction at a <sup>7</sup>Li beam energy of 32 MeV...

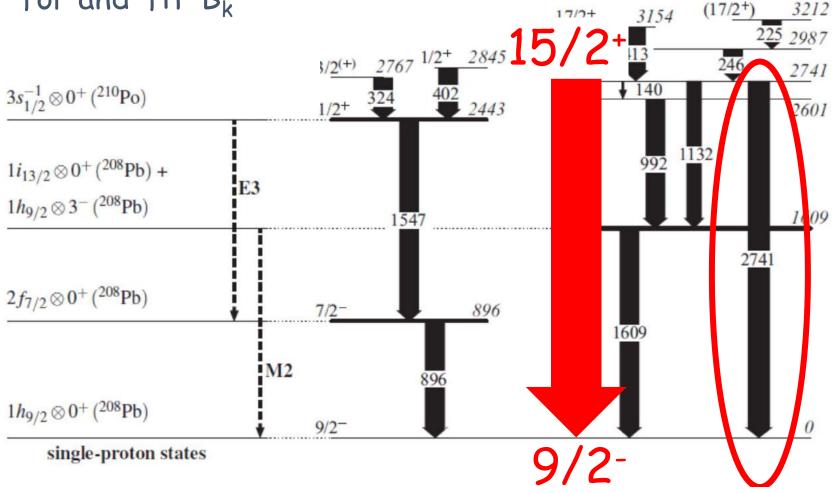
...it's a sort of compound nucleus reaction but

<sup>7</sup>Li breaks up into <sup>4</sup>He and <sup>3</sup>H so is the <sup>4</sup>He just a

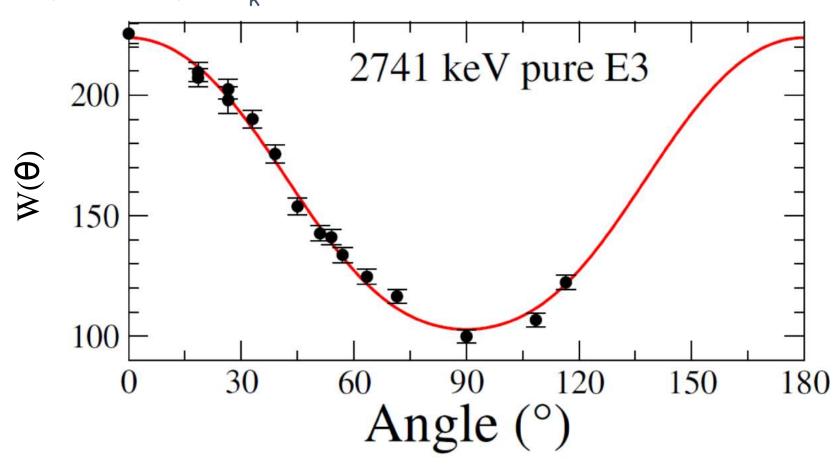
spectator?

Anyway, the question is how much alignment is in the initial states in <sup>209</sup>Bi??

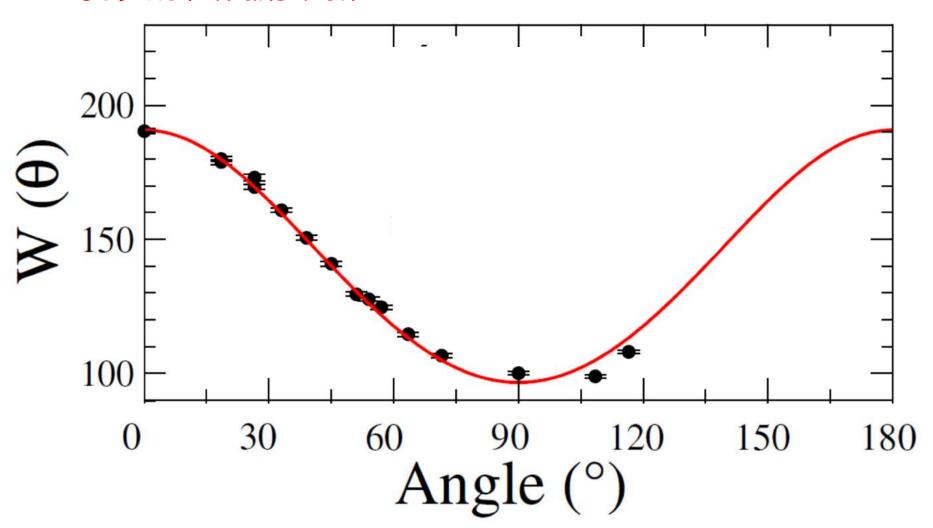
Answer...look at something we know the  $A_k$  values for and 'fit'  $B_k$ 



Answer...look at something we know the  $A_k$  values for and 'fit'  $B_k$ 

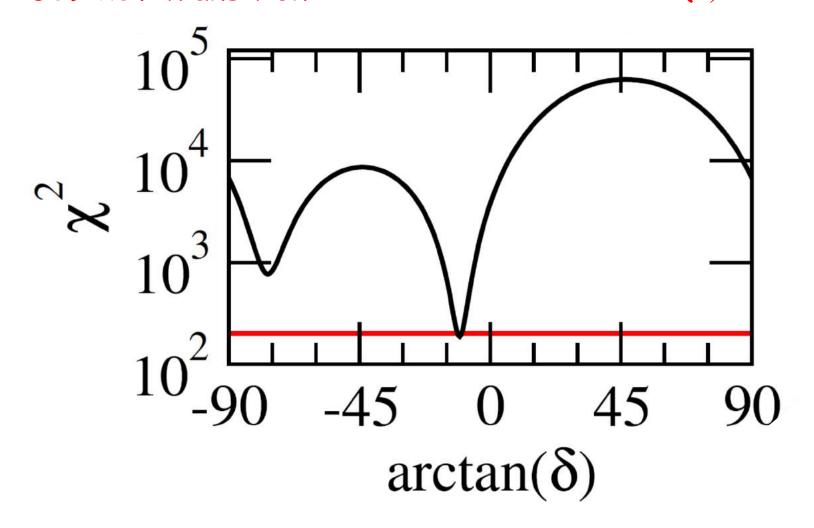


#### 1609 keV transition



1609 keV transition

min at  $arctan(\delta) = -10.54$ 



O.J.Roberts, C.R.Nita et al., Phys Rev C93 (2016) 014309

#### 1609 keV transition:

$$arctan(\delta) = -10.54$$

$$\delta$$
 (E3/M2) = -0.184(13)

%E3 = 
$$\frac{\delta^2}{1+\delta^2}$$
 = 3%

%M2 = 
$$\frac{1}{1+\delta^2}$$
 = 97%

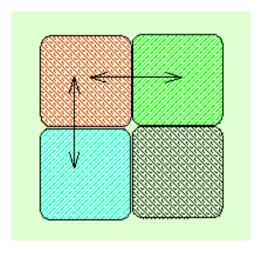
## What's the physics?

$$\frac{13^{+}}{2} = \frac{\left(0^{+} \otimes \pi 1 i_{13/2}\right)}{8} 1609$$

$$\frac{9^{-}}{2} = \frac{9^{-}}{\left(0^{+} \otimes \pi 1 h_{9/2}\right)} 0$$

O.J.Roberts, C.R.Nita et al., Phys Rev C93 (2016) 014309

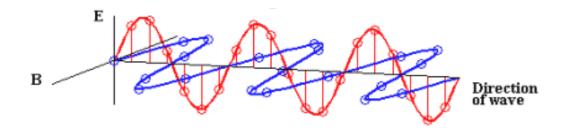
# Linear polarisation



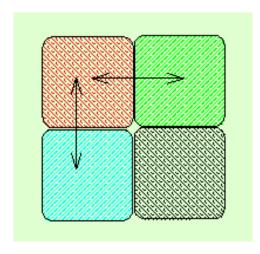
γ ray travelling into the slide

A segmented detector can be used to measure the linear polarisation which can be used to distinguish between magnetic (M) and electric (E) character of radiation of the same multipolarity.

The Compton scattering cross section is larger in the direction perpendicular to the electric field vector of the radiation.



# Linear polarisation



γ ray travelling into the slide

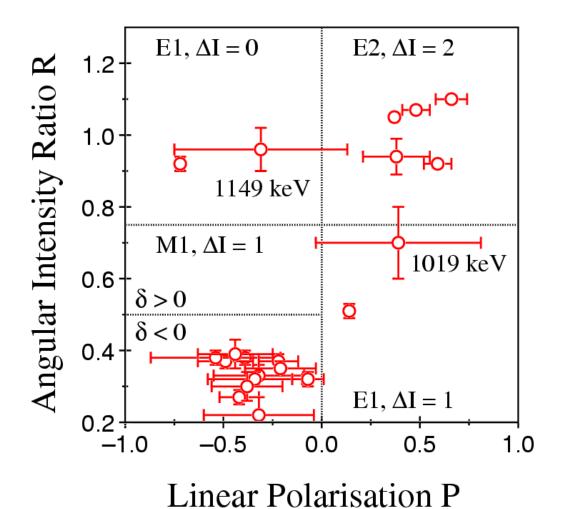
Define experimental asymmetry as:

$$A = \frac{N_{90} - N_0}{N_{90} + N_0}$$

where  $N_{90}$  and  $N_0$  are the intensities of scattered photons perpendicular and parallel to the reaction plane.

The experimental linear polarisation P = A/Q where Q is the polarisation sensitivity of the dectector.

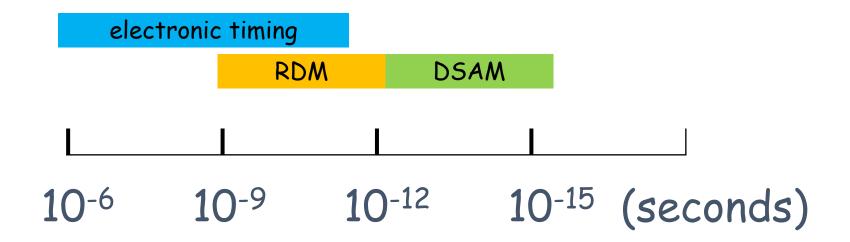
## Linear polarisation



Plot P against the angular distribution information to uniquely define the multipolarity.

Data from Eurogam

# Measuring level lifetimes



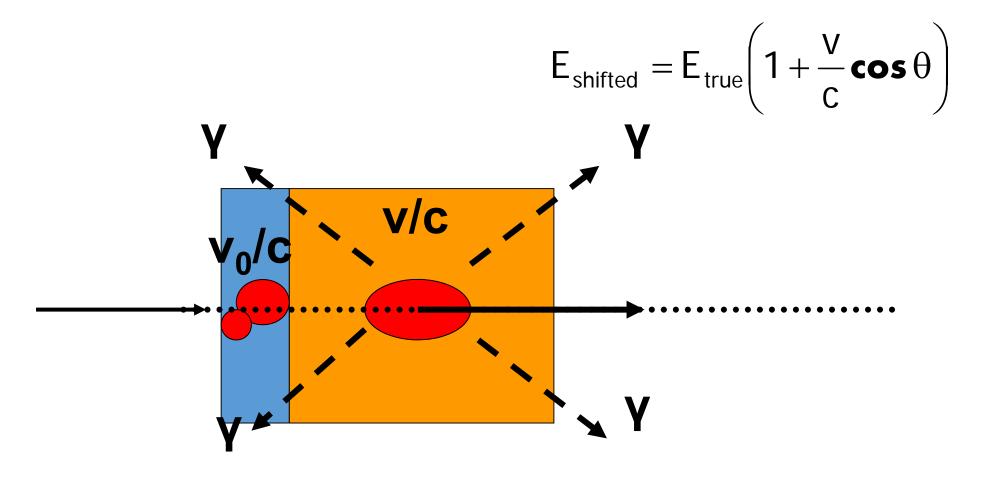
DSAM - Doppler shift attenuation method

RDM - Recoil distance method

Electronic timing - Using arrays of fast scintillation detectors

## Doppler shift attenuation method

- Measure lifetimes in the range  $10^{-15} < \tau < 10^{-12}$  s.
- Stopping time in metal foil is comparable to lifetimes of excited states.



#### Centroid shift method

$$E_{f} = E_{true} \left( 1 + \frac{V}{C} \cos \theta_{f} \right)$$

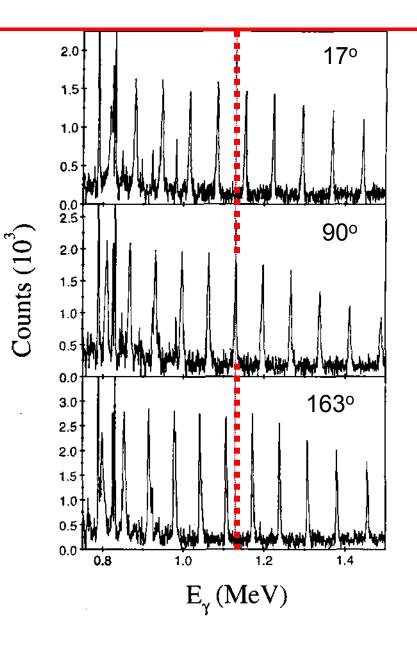
$$E_{b} = E_{true} \left( 1 + \frac{V}{C} \cos \theta_{b} \right)$$

If 
$$\theta = \theta_{forward} = 180 - \theta_{backward}$$

$$E_f - E_b = 2E_{true} \frac{V}{c} \cos \theta$$

$$F(\tau) = \frac{V}{V_0}$$
 and  $\beta = \frac{V_0}{C}$ 

$$F(\tau) = \frac{E_f - E_b}{2E_{true}\beta \cos \theta}$$



#### Recoil distance method

Have a space between the target and the stopper.

Measuring the v/c you can work out whether the gammaray is emitted before/after the stopper

Vary the target/stopper distance to get a measurement of the lifetime of the level

### Recoil distance method

$$I_{degraded} = Ie^{-d/v\tau}$$

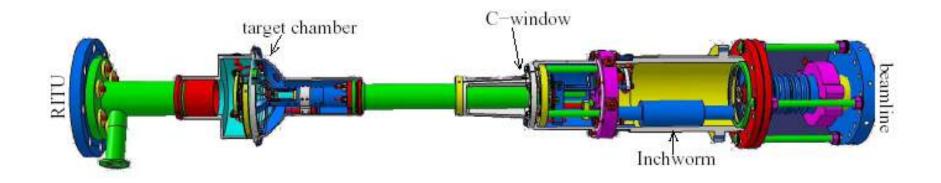
$$I_{shifted} = I(1 - e^{-d/v\tau})$$

$$\frac{I_{degraded}}{I_{degraded} + I_{shifted}} = e^{-d/v\tau}$$

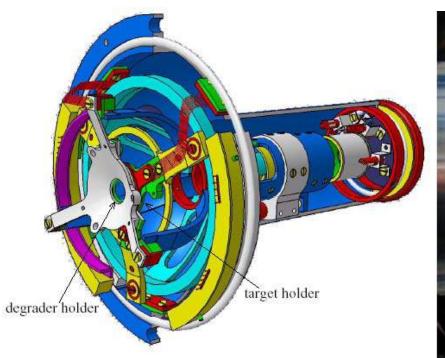
$$\frac{I_{degraded} + I_{shifted}}{I_{degraded}} = e^{-d/v\tau}$$

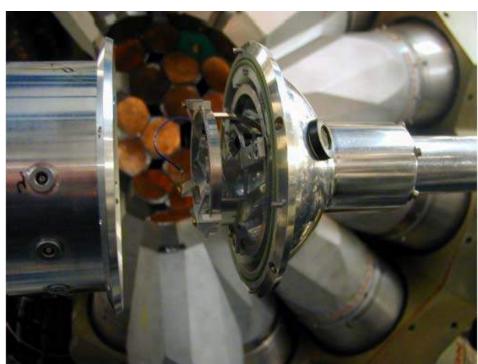
$$\frac{I_{degraded}}{I_{degraded}} = e^{-d/v\tau}$$

$$\frac{I_{degraded}}{I_{$$

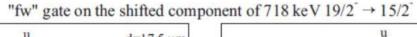


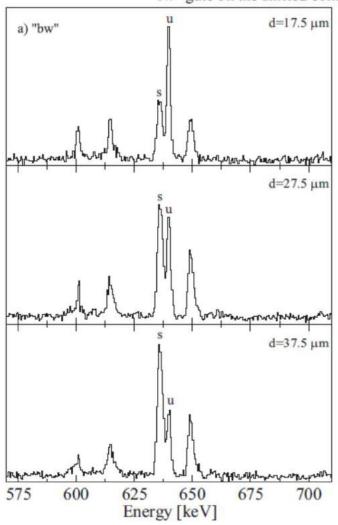
#### **The Koln Plunger Device**

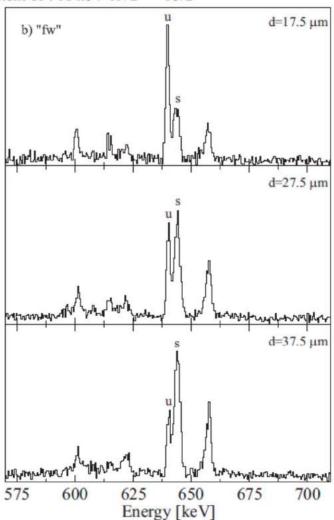




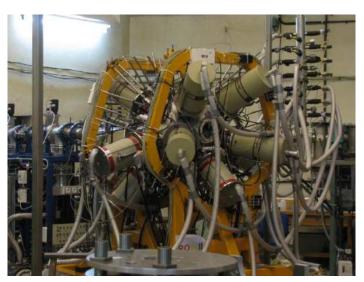
#### Lifetime of the 15/2-state in <sup>119</sup>Te







# Direct measurement using ROSPHERE at Bucharest:



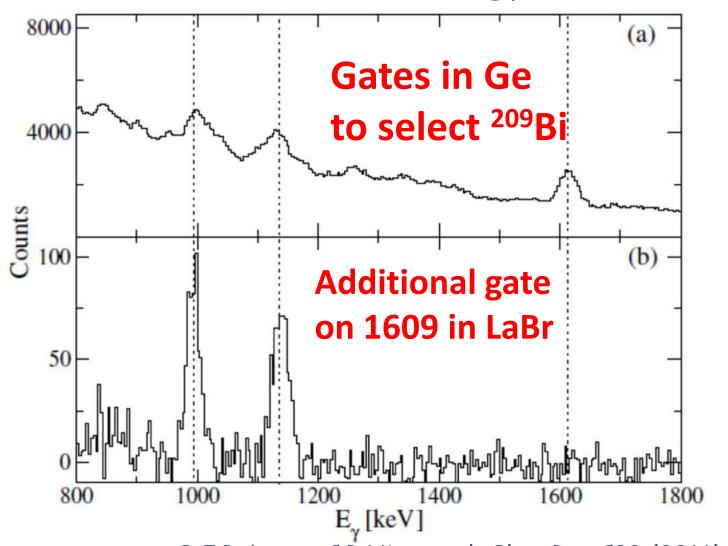
- 15 HPGe detectors (A/C):
  - 10 x HPGe detectors @ 37°
  - 1 x HPGe detector @ 64°
  - 4 x HPGe detectors @ 90°



- 11 LaBr<sub>3</sub>(Ce):
  - $\emptyset2"x2"$  @ 90 and 64° (three) (Cylindrical)
  - ø1.5"x1.5" @ 90 (six)(Cylindrical)
  - ø1"x1.5" @ 64° (two) (Conical)

## ROSPHERE at Bucharest: example of 209Bi

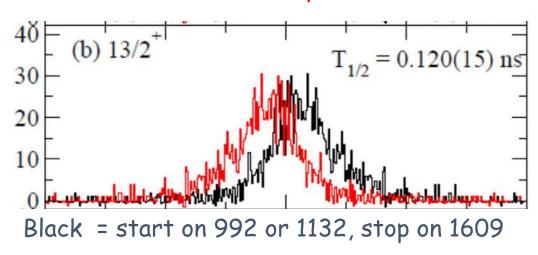
 $^{208}$ Pb( $^{7}$ Li, $^{209}$ Bi at beam energy of 32 MeV

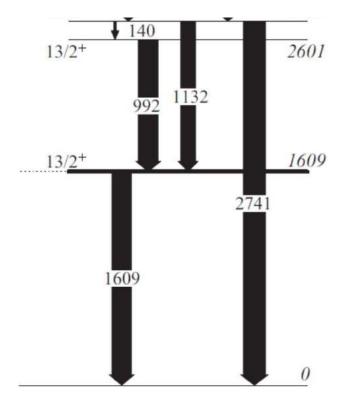


O.J.Roberts, C.R.Nita et al., Phys Rev C93 (2016) 014309

## ROSPHERE at Bucharest: example of 209Bi







Difference in the centroids is  $2\tau$  (give or take some correction factors)

O.J.Roberts, C.R.Nita et al., Phys Rev C93 (2016) 014309

## What's the physics?

$$\frac{13^{+}}{2} = \frac{\mathsf{t}_{1/2} = 120\mathsf{ps}}{\mathsf{E3}} = \frac{\left(0^{+} \otimes \pi I i_{13/2}\right)}{\mathsf{M2} < 1 \; \mathsf{Wu}} = \frac{9^{-}}{2} = \frac{\left(0^{+} \otimes \pi I h_{9/2}\right)}{\left(0^{+} \otimes \pi I h_{9/2}\right)} = \frac{\left(0^{+} \otimes \pi I h_{9/2}\right)}{\left(0^{+} \otimes \pi I h_{9/2}\right)} = \frac{\mathsf{M2} + \mathsf{M2} + \mathsf{M3} + \mathsf{M3}$$

O.J.Roberts, C.R.Nita et al., Phys Rev C93 (2016) 014309

### From first lecture:

Typical transition rates:

$$T(E1) = 1.59 \times 10^{15} (E_{\gamma})^3 B(E1)$$

$$T(E2) = 1.22 \times 10^9 (E_y)^5 B(E2)$$

$$T(M1) = 1.76 \times 10^{13} (E_{\gamma})^3 B(M1)$$

$$T(M2) = 1.35 \times 10^7 (E_y)^5 B(M2)$$

 $E_{\gamma}$  in MeV

B(EI) in  $e^2fm^{2\lambda}$ 

B(MI) in 
$$\left(\frac{e\hbar}{2Mc}\right)^2$$
 fm<sup>2 $\lambda$ -2</sup>

$$\frac{1}{\tau} = T (\sigma \lambda)$$

If E2:

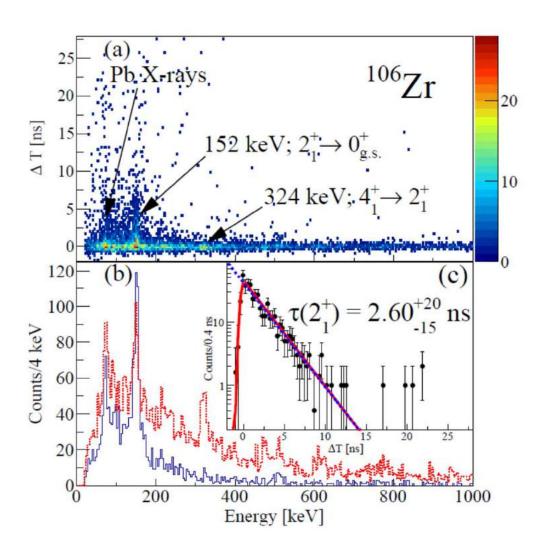
$$T(E2) = 1.22 \times 10^9 (E_y)^5 B(E2)$$

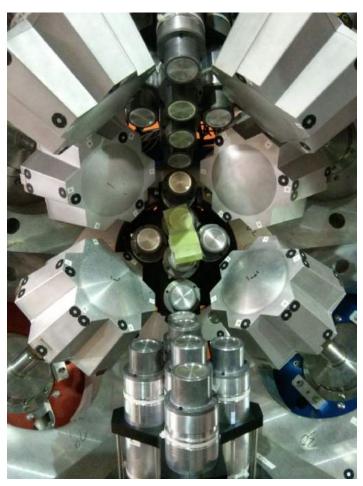
The transition quadrupole moment  $Q_0$  is obtained from:

$$Q_0^2 = \frac{16\pi B(E2)}{5 \langle J_i K20 | J_f K \rangle^2}$$

So measuring a lifetime gives us a transition quadrupole moment (note that we cannot get the sign of  $Q_0$ )

## Using the EURICA/FATIMA array at RIKEN





Browne et al. PLB750 (2015) 448

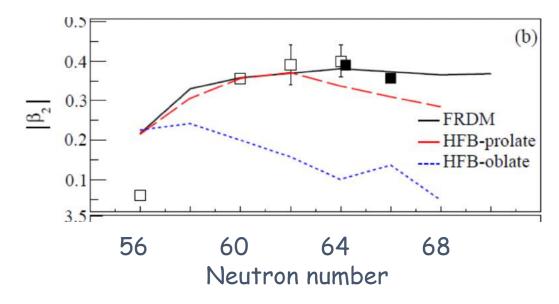
## Quadrupole moment relates to shape

Assuming a quadrupoloid shape and that the deformation is the same

for both states, the deformation  $\beta_2$  can be obtained from:

$$Q_0 = \frac{3}{\sqrt{5\pi}} ZR^2 \beta_2 \left( 1 + \frac{2}{7} \sqrt{\frac{5}{\pi}} \beta_2 + \frac{1}{14\pi} \beta_2^2 + \cdots \right)$$

#### Neutron-rich Zr nuclei



Formula: Löbner, Vetter and Honig, Nucl. Data Tab A7 (1970) 495.

Data: Browne et al. PLB750 (2015) 448

# Suggestions for tutorial discussion:

- 1. In the angular correlation formula, on what might  $Q_k$  depend?
- 2. How does the deformation parameter  $\beta_2$  relate to  $\epsilon_2$  and  $\delta$  (which you may see elsewhere)
- 3. What methods are there to measure the sign of the quadrupole moment?



Acknowledgements (for slides and animations)

Dr Dave Joss

Prof John Simpson

# Useful points of reference for angular correlation and distribution

Chapters 12,14 and 15 in 'The electromagnetic interaction in nuclear spectroscopy' edited by W.D.Hamilton

Q factors: Camp and van Lehn, NIM 76 (1969) 192

Alignment in compound nuclear reactions:
Butler and Twin, NIM 190 (1981) 283

Errors on  $\delta$  from arctan  $\delta$  plots: James, Twin and Butler, NIM 115 (1974) 105