

### Experimental Nuclear Structure Part I

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Argonne National Laboratory



A U.S. Department of Energy Office of Science Laboratory Operated by The University of Chicago





## Outline

- I) Lecture I: Experimental nuclear structure physics
  - reactions used to populate excited nuclear states
  - □ techniques used to measure the lifetime of a nuclear state
    - Coulomb excitation, electronic, specific activity, indirect
  - techniques used to deduce J<sup>π</sup>
    - > ICC, angular distributions, DCO ratios etc.
- II) Lecture II: Contemporary Nuclear Structure Physics at the Extreme
  - spectroscopy of nuclear K-Isomers
  - **physics** with large γ-ray arrays
  - **gamma-ray tracking** the future of the γ–ray spectroscopy

Have attempted to avoid formulas and jargon, and material covered by other lecturers – will give many examples

Please feel free to interrupt at any time!





## Some Useful Books

- "Handbook of nuclear spectroscopy", J. Kantele,1995
- "Radiation detection and measurements", G.F. Knoll, 1989
- "In-beam gamma-ray spectroscopy", H. Morinaga and T. Yamazaki, 1976
- "Gamma-ray and electron spectroscopy in Nuclear Physics", H. Ejiri and M.J.A. de Voigt, 1989
- "Techniques in Nuclear Structure Physics", J.B.A. England, 1964
- "Techniques for Nuclear and Particle Physics Experiments", W.R. Leo, 1987
- "Nuclear Spectroscopy and Reactions", Ed. J. Cerny, Vol. A-C
- "Alpha-, Beta- and Gamma-ray Spectroscopy", Ed. K. Siegbahn, 1965
- "The Electromagnetic Interaction in Nuclear Spectroscopy", Ed. W.D. Hamilton, 1975

Plenty of information on the Web







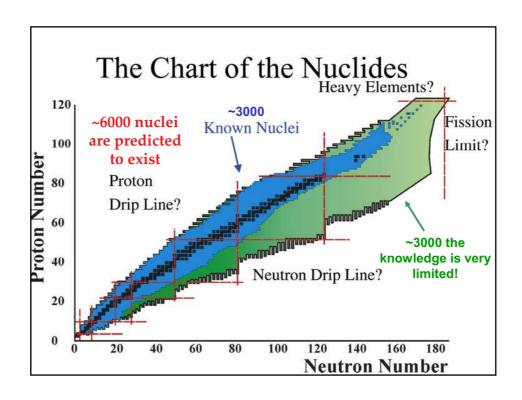
# Input from many colleagues

- C.J. Lister and I. Ahmad, Argonne National Laboratory, USA
- M.A. Riley, Florida State University, USA
- I.Y. Lee, Lawrence Berkeley National Laboratory, USA
- D. Radford, Oak Ridge National Laboratory, USA
- A. Heinz, Yale University, USA
- C. Svensson, University of Guelph, Canada
- G.D. Dracoulis and T. Kibedi, Australian National University, Australia
- J. Simpson, Daresbury Laboratory, UK
- E. Paul, University of Liverpool, UK
- P. Reagan, University of Surrey, UK

and many others ...







## Introduction

- ☐ The nucleus is one of nature's most interesting quantal few-body systems
- ☐ It brings together many types of behaviour, almost all of which are found in other systems
- ☐ The major elementary excitations in nuclei can be associated with single-particle and collective modes.
- ☐ While these modes can exist in isolation, it is the interaction between them that gives nuclear spectroscopy its rich diversity





### So to summarize ...

### NUCLEAR PHYSICS IS A BIG CHALLENGE

(because of complicated forces, energy scale, and sizes involved)

The challenge of understanding how nucleon-nucleon interactions build to create the mean field or how single-particle motions build collective effects like pairing, vibrations and shapes

#### NUCLEAR PHYSICS IS IMPORTANT

(intellectually, astrophysics, energy production, and security)

#### THIS IS A GREAT TIME IN NUCLEAR PHYSICS

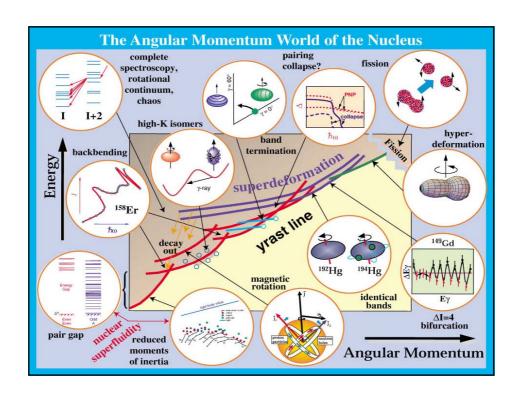
(with new facilities just around the corner we have a chance to make major contributions to the knowledge - with advances in theory we have a great chance to understand it all - by compiling & evaluating data we have a chance to support various applications and to preserve the knowledge for future generations!)

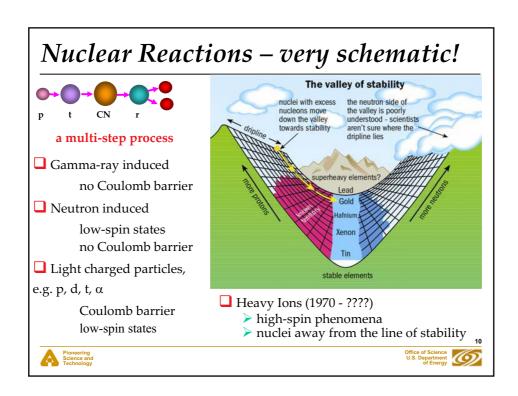


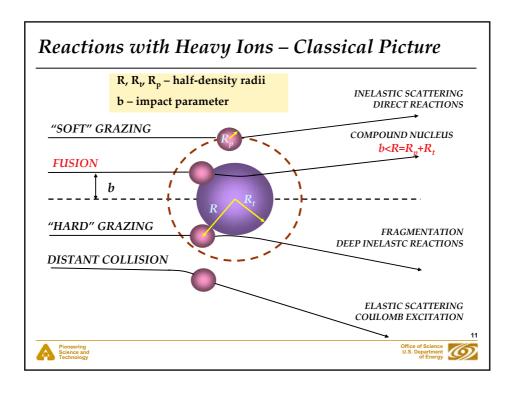
Large N/Z To learn many of the secrets of the nucleus -

> we have to put it at extreme conditions and study how it survives

such a stress!







## Heavy Ions at the Coulomb barrier

Many properties of the collision can be quite well estimated by just using conservation of momentum and energy.

$$E_{cm} = M_t / (M_b + M_t) E_{lab}$$

Energy scale on which fusion starts is determined by Coulomb barrier,  $V_{\rm ch}$ 

$$V_{cb} = (4\pi\epsilon)^{-1} Z_b Z_t e^2 / R = 1.44 Z_b Z_t / 1.16 [(A_b^{-1/3} + A_t^{-1/3}) + 2] MeV$$

$$L_{max} = 0.22 \; R \; [\; \mu \; (E_{cm} - V_{cb}) \;]^{1/2} \qquad \hbar$$

Excitation energy is usually lowered by Q-value and K.E. of evaporated particles

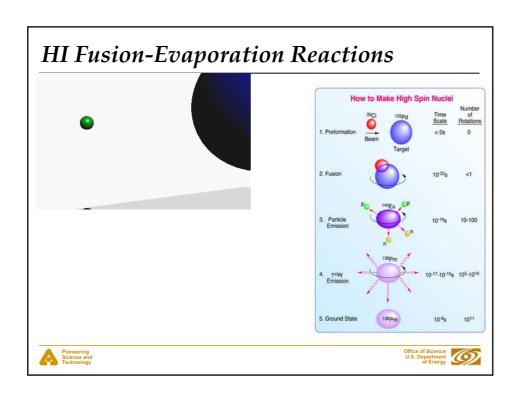
$$E_{residue}^* = E_{cm} + Q - K.E.$$

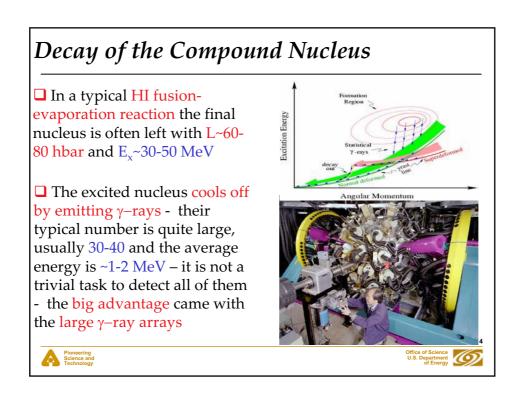
Velocity of center-of-mass frame, which is ~ velocity of fused residues

$$\beta_{\rm r}^2 = 2 M_b c^2 E_{\rm lab} / [(M_b + M_t)c^2]^2$$









## Channel Selection for γ–ray spectroscopy

#### Detection of Light Charged Particles (α,p,n)

PLUS Efficient, flexible, powerful....inexpensive.

MINUS Count-rate limited, Contaminant (Carbon etc, isotopic impurities) makes absolute identification of new nuclei difficult.

that is,  $\sim 10^{-4}$ CROSS SECTION LOWER LIMIT ~100 μb

Detection of Residues in Vacuum Mass Separator

PLUS True M/q, even true M measurement. With suitable focal plane detector can be ULTRA sensitive. Suppresses contaminants.

MINUS Low Efficiency

that is  $\sim 10^{-7}$ CROSS SECTION LOWER LIMIT ~100 nb

Detection of Residues in Gas Filled Separator

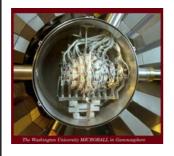
Improves efficiency of vacuum separators, at cost of mass information and clenliness. In some cases (heavy nuclei) focal plane counters clean up the data for good sensitivity.







## Some Channel Selection Detectors



Argonne FMA USA



Light charged-particle detector Microball - 96 CsI with photo diodes USA



Jyvaskyla **RITU** Europe



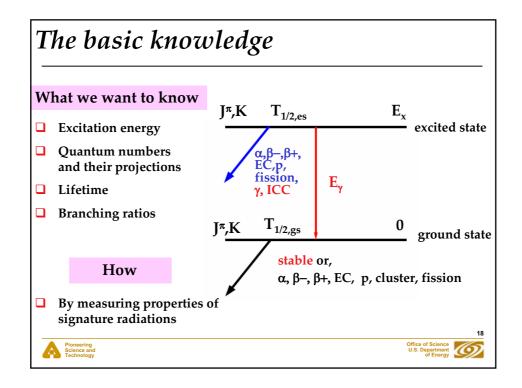


Reaction Yield/sec.  $Y = N_b \ N_t \ \sigma$  with  $i_b$  = electric current in amps, q = charge state, e = 1.6 10-19 c  $N_t = [N_a / A] \ \rho x$  with  $N_a$  = Avagadros #, A = Mass # of tgt,  $\rho$  = density in g/cc, x = thickness (cm).  $\sigma$  = Cross Section in cm² .... note 1 barn is  $10^{-24}$  cm

Accumulated data:  $D = Y \ x \ TIME \ x \ Efficiency$ Typical "far from stability" near barrier experiment may have:  $i_b = 100 \ nA$ , q=10, A=100,  $\rho x$ =10-3 &  $\sigma$ =1 barn - produces 3x10-5 reactions/sec

BUT

If partial cross section is 100 nb and efficiency is 10%..... rate is 10 /hour, 10 pb gives ~ 1 every 10 weeks!!!.....the present situation for producing the heavies elements.



### What is Stable?

A surprisingly difficult question with a somewhat arbitrary answer! CAN'T Decay to something else, BUT

#### CAN'T Decay is a Philosophical Issue

☐ Violation of some quantity which **we believe** is conserved such as Energy, Spin, Parity, Charge, Baryon or Lepton number, etc.

### DOESN'T Decay is an Experimental Issue that backs up the beliefs

Specific Activity:  $A=dN/dt = \lambda N$ 

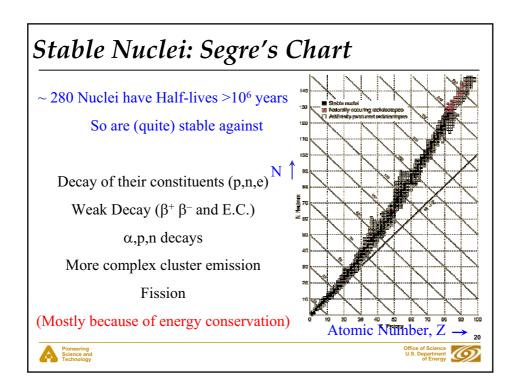
Activity of 1 mole of material (6.02 x  $10^{23}$  atoms) with  $T_{1/2}=10^9$  v ( $\lambda=2.2$  x  $10^{-17}$  s) is ~0.4 mCi (1 Ci=3.7 x  $10^{10}$  dps) (or 13 MBq) .... a blazing source, so it is quite easy to set VERY long limits on stability.

Current limit on proton half-life, based on just counting a tank of water is  $T_{1/2} > 1.5 \times 10^{25} \text{ yr.}$ 









# Mean Lifetime

$$f_{decay}(t) = \frac{Ae^{-\lambda t}}{\int\limits_{0}^{\infty} Ae^{-\lambda t} dt} = \lambda e^{-\lambda t}$$

Probability for decay of a nuclear state (normalized distribution function);  $\lambda$  – decay constant

$$1 - P_n(t) = 1 - \int_0^t \lambda e^{-\lambda t'} dt' = e^{-\lambda t}$$

Probability that a nucleus will remain at time t

$$P_n(t) = \int_0^t \lambda e^{-\lambda t'} dt'$$

Probability that a nucleus will decay within time t

$$\langle t \rangle = \tau = \int_{0}^{\infty} t e^{-\lambda t} dt = \frac{1}{\lambda}$$

The average survival time (mean lifetime -  $\tau$ ) is then the mean value of this probability

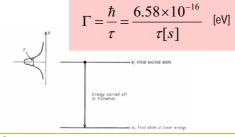


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# Half-life & Decay Width

 $T_{1/2}$ : the time required for half the atoms in a radioactive substance to disintegrate

relation between  $\tau$ ,  $T_{1/2}$  and  $\lambda$   $\tau = \frac{T_{1/2}}{\ln 2} = \frac{1}{\lambda}$ 

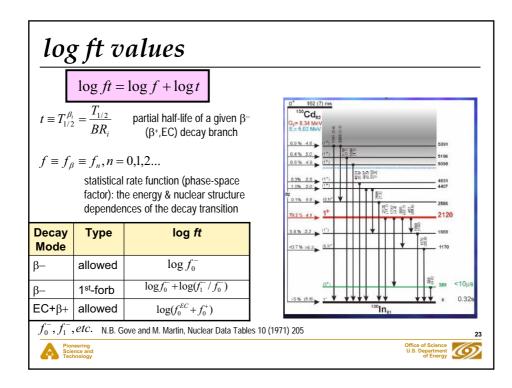


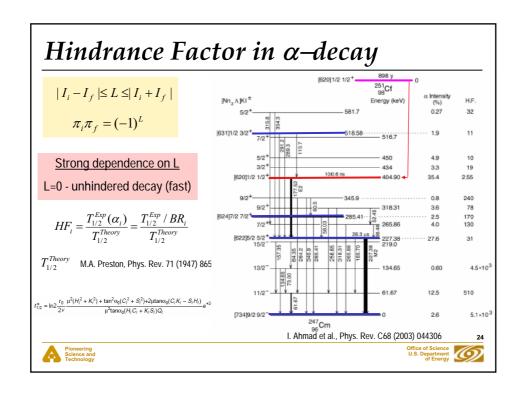
 $A = A_0 2^{\frac{1}{1/12}}$   $= A_0 \theta^{\frac{1}{1/12}}$   $= A_0 \theta^{\frac{1}{1/12}}$   $= A_0 \theta^{\frac{1}{1/12}}$   $= A_0 \theta^{\frac{1}{1/12}}$   $= A_0 \theta^{\frac{1}{1/12}}$  0 T 2T 3T 4T 5TTime as a multiple of the halfille T

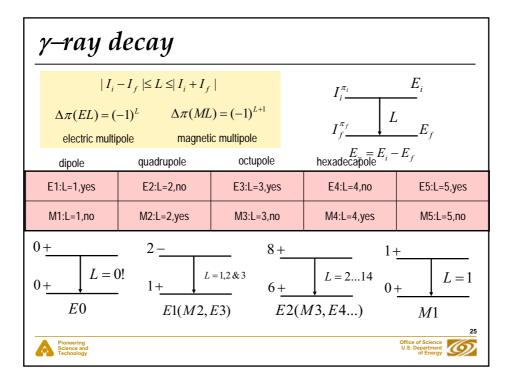
 $\Gamma \propto |\langle \psi_1 | M | \psi_2 \rangle|^2$ 

Determine the matrix element describing the mode of decay between the initial and final state

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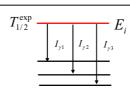






# Partial lifetime & Transition Probability

$$T_{1/2}^{\gamma} = T_{1/2}^{\exp} \times \frac{\sum I_{\gamma_i} \times (1 + \alpha_{T_i})}{I_{\gamma}}$$
 partial half-life



$$P_{\gamma}(XL:I_{i}\to I_{f}) = \frac{\ln 2}{T_{1/2}^{\gamma}} = \frac{8\pi(L+1)}{L[(2L+1)!!]^{2}} \left(\frac{E_{\gamma}}{\hbar c}\right)^{2L+1} B(XL:I_{i}\to I_{f})$$

<u>partial γ–ray</u> Transition Probability

reduced Transition Probability

$$B(XL:I_i \to I_f) = \frac{\left| \left\langle I_i \middle| M(XL) \middle| I_f \right\rangle \right|^2}{2I_i + 1}$$

contains the nuclear structure information





# Hindrance Factor in $\gamma$ -ray decay

 $F_{W(N)} = \frac{B(XL)_{Theory}}{B(XL)_{Exp}} = \frac{T_{1/2}^{\gamma}(XL)_{Exp}}{T_{1/2}^{\gamma}(XL)_{Theory}}$ 

<u>Hindrance Factor: Weisskopf (W):</u> based on spherical shell model potential

> Nilsson (N): based on deformed Nilsson model potential

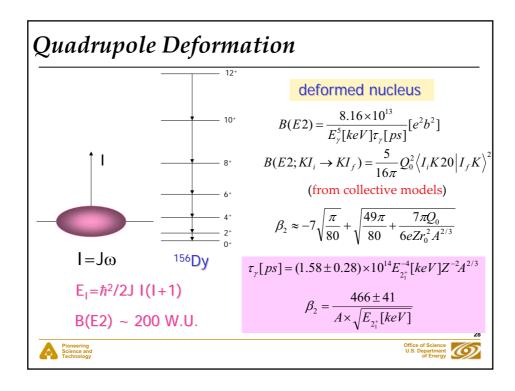
... usually an upper limit, but ...

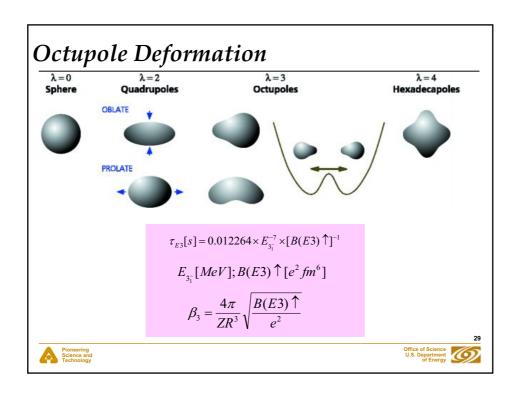
EL	$B(EL)_W, e^2 fm^{2L}$	$T_{1/2}^{\gamma}(EL)_{W}$ , sec	ML	$B(ML)_{W}, \mu_{N}^{2} fm^{2L-2}$	$T_{1/2}^{\gamma}(ML)_{W}$ , sec
E1	$0.06446A^{2/3}$	$6.762A^{-2/3}E_{\gamma}^{-3}\times10^{-15}$	M1	1.7905	$2.202E_{\gamma}^{-3} \times 10^{-14}$
E2	$0.0594A^{4/3}$	$9.523A^{-4/3}E_{\gamma}^{-5}\times10^{-9}$	M2	$1.6501A^{2/3}$	$3.100A^{-2/3}E_{\gamma}^{-5}\times10^{-8}$
E3	$0.0594A^2$	$2.044A^{-2}E_{\gamma}^{-7}\times10^{-2}$	М3	$1.6501A^{4/3}$	$6.655A^{-4/3}E_{\gamma}^{-7}\times10^{-2}$
E4	$0.06285A^{8/3}$	$6.499A^{-8/3}E_{\gamma}^{-9}\times10^{4}$	M4	$1.7458A^2$	$2.116A^{-2}E_{\gamma}^{-9}\times10^{5}$
E5	$0.06929A^{10/3}$	$2.893A^{-10/3}E_{\gamma}^{-11}\times10^{11}$	M5	$1.9247A^{8/3}$	$9.419A^{-8/3}E_{\gamma}^{-11}\times10^{11}$

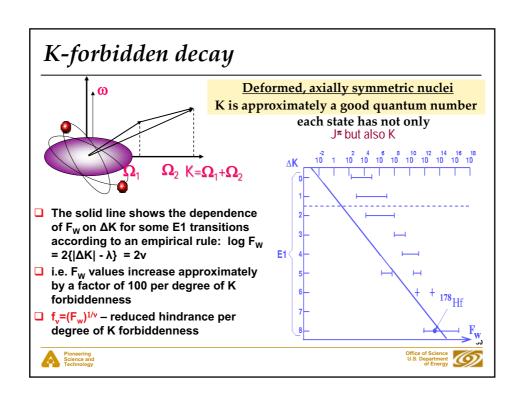


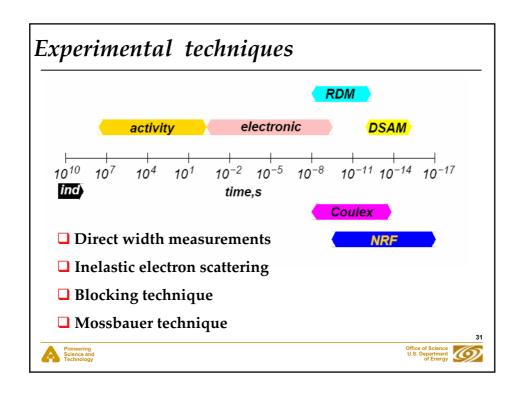


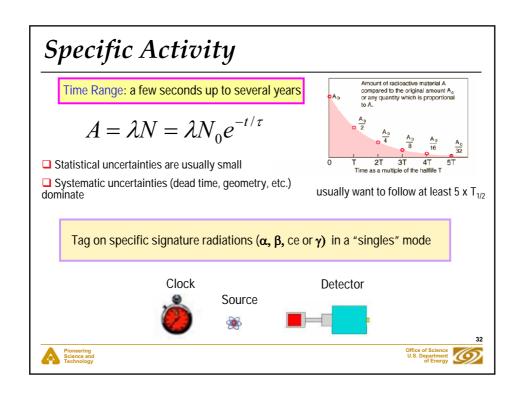


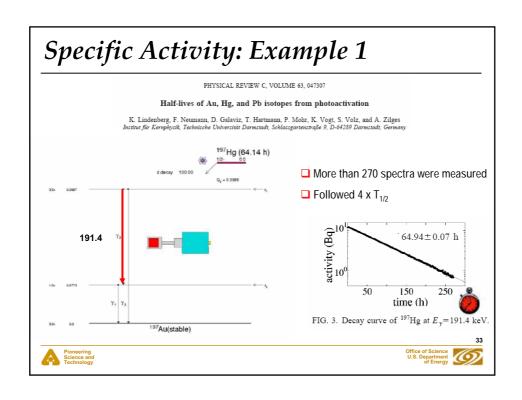


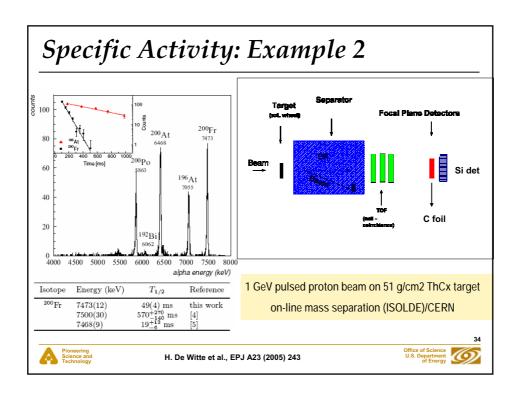


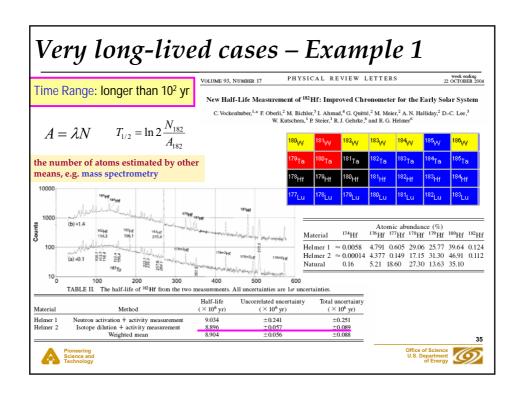


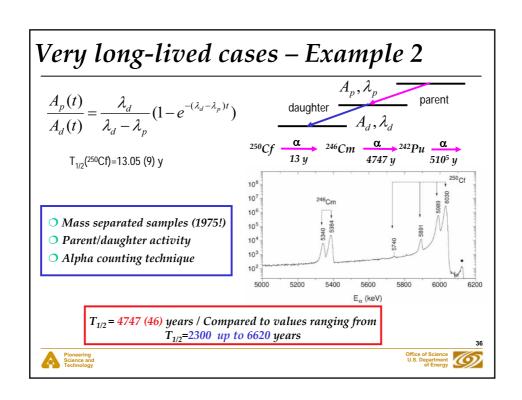


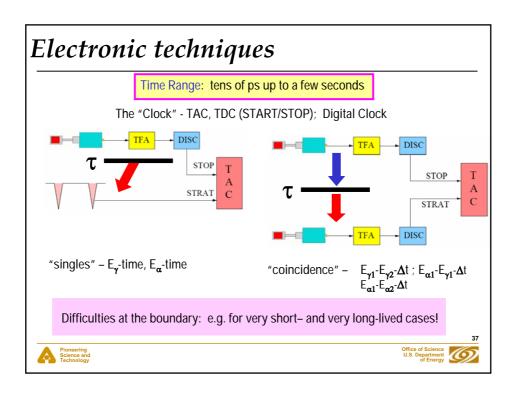


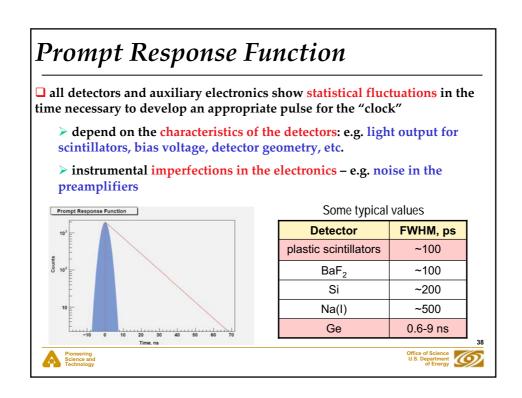


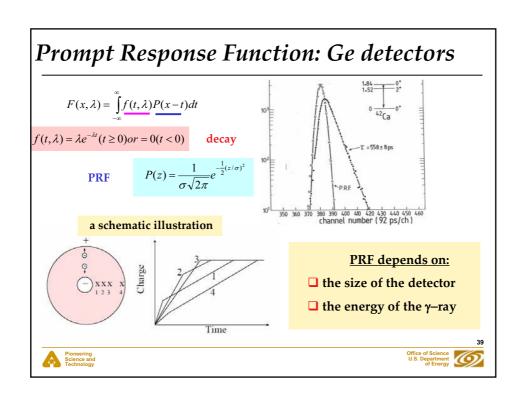


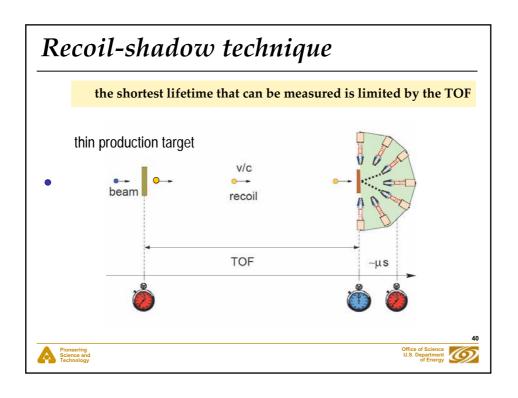


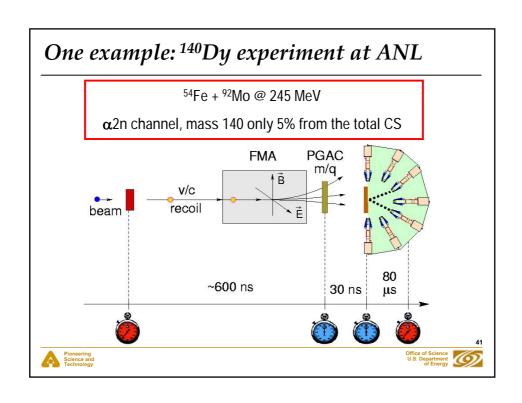


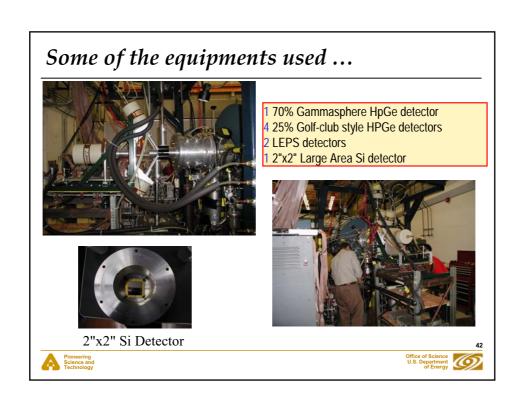


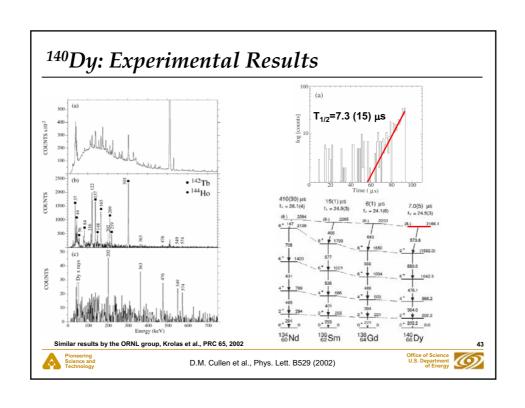


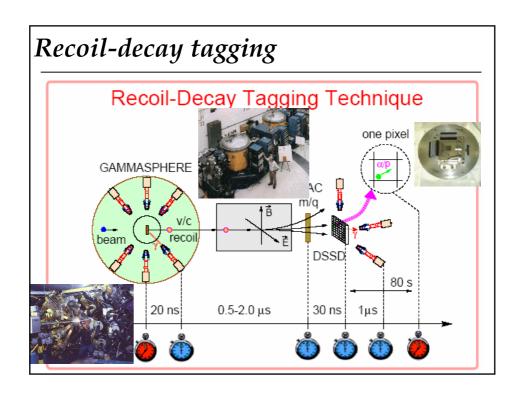


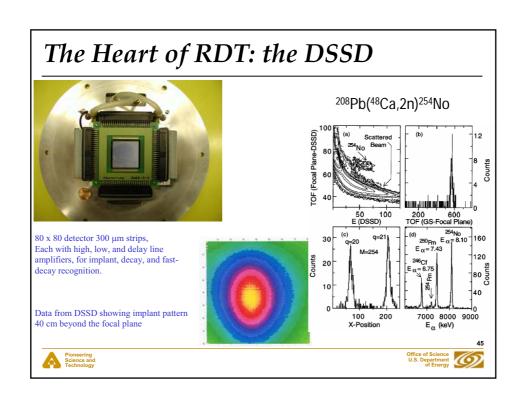


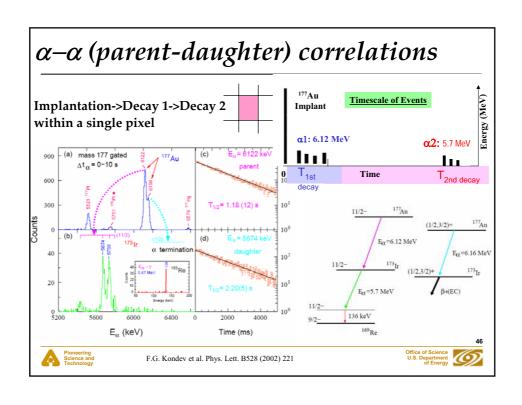


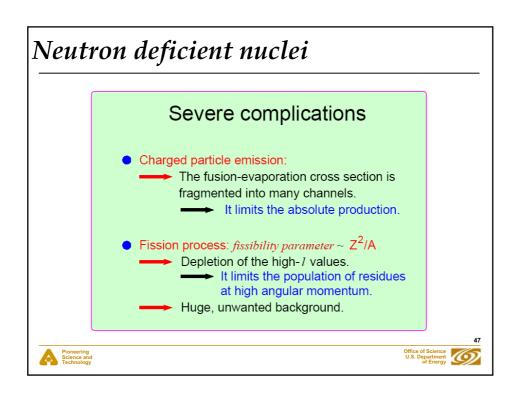


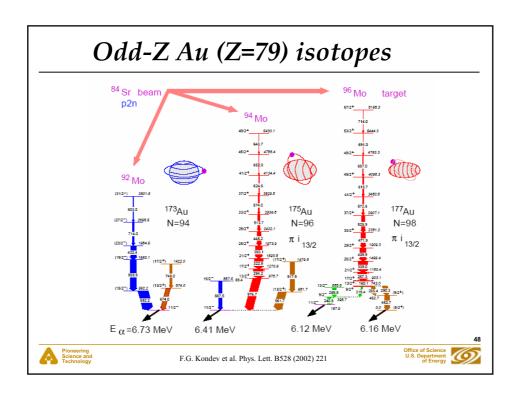


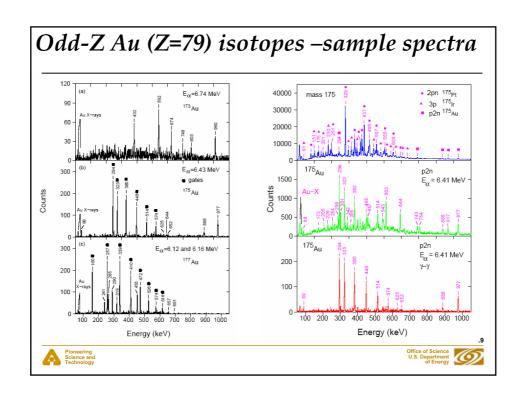


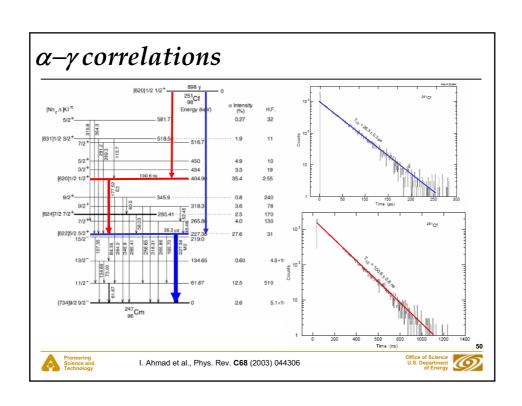


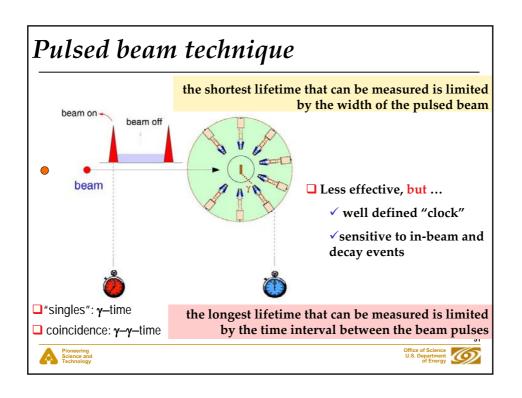


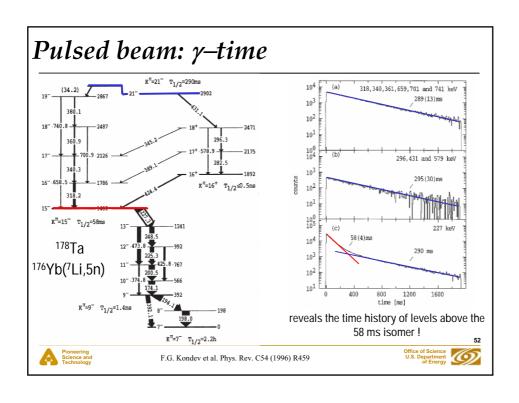


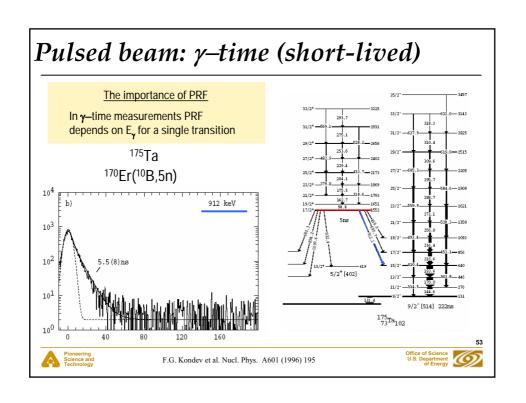


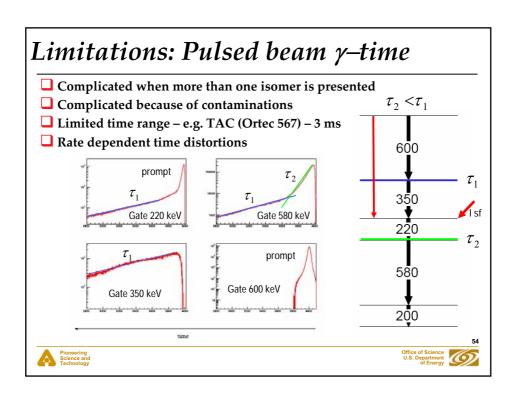


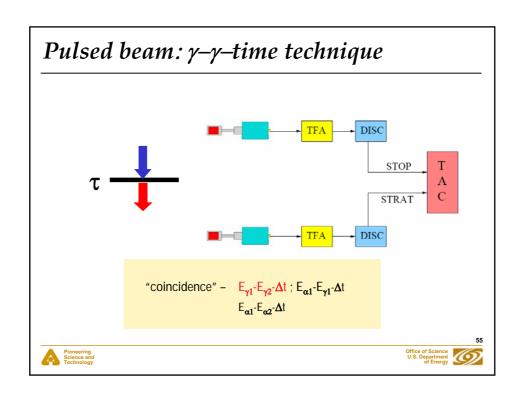


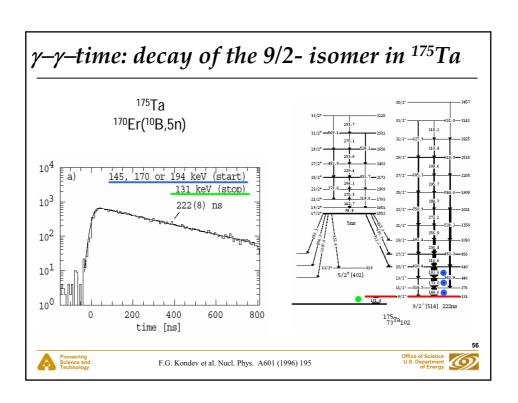


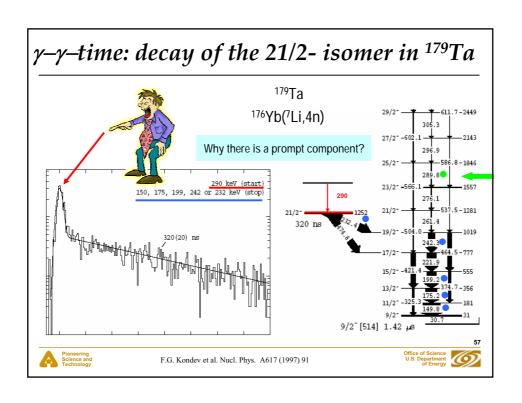


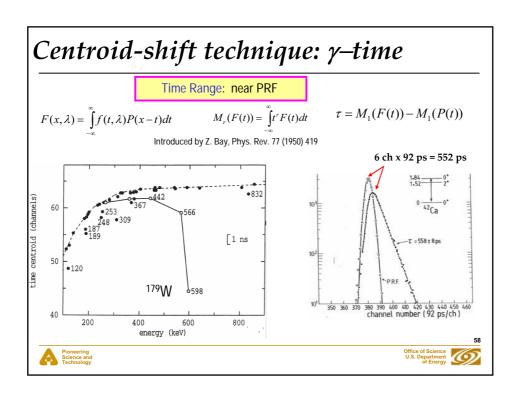


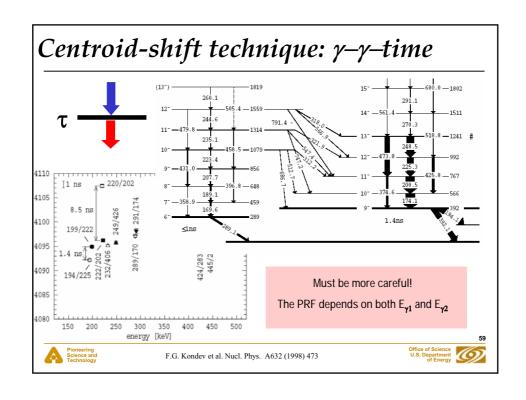


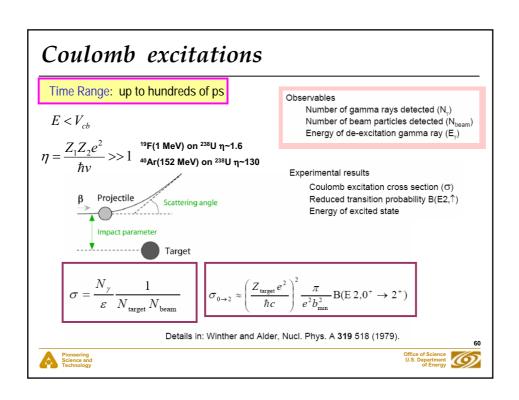


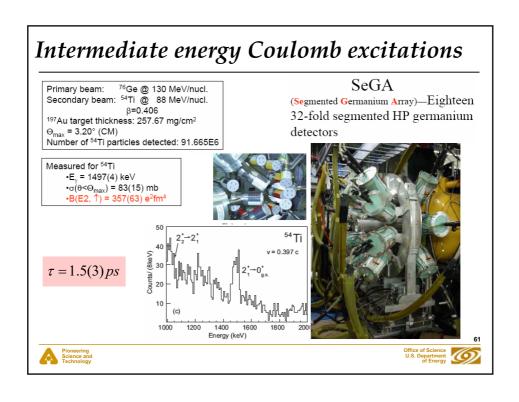


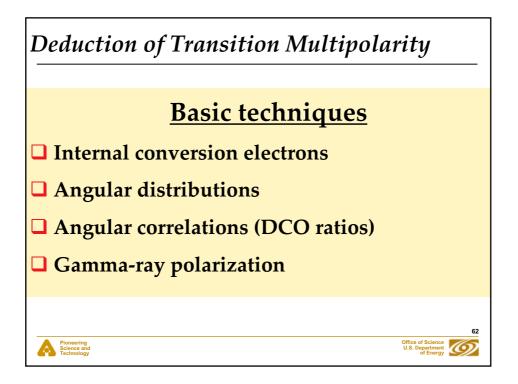


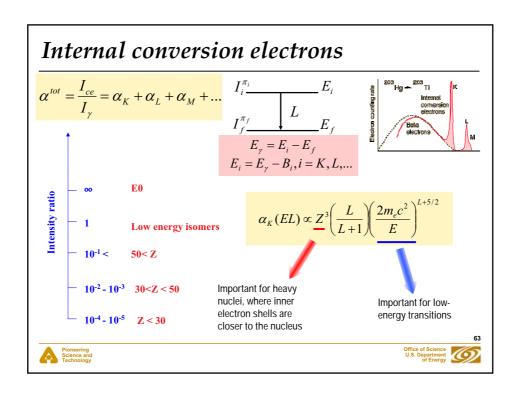


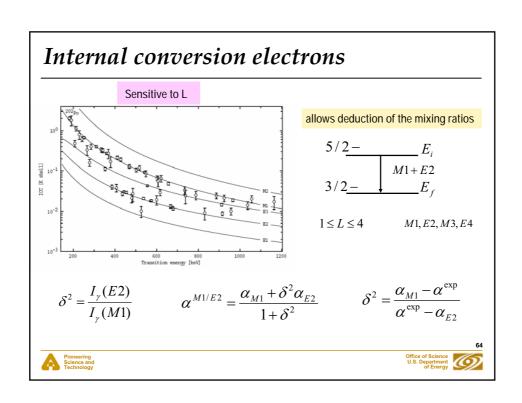


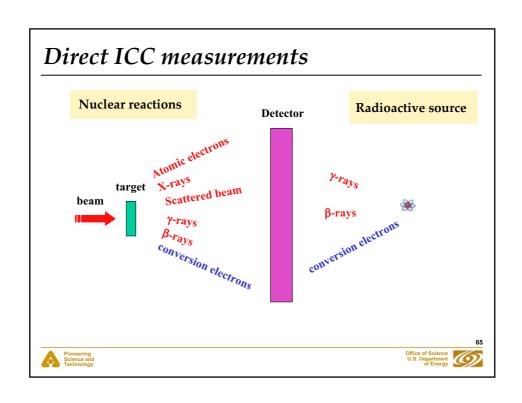


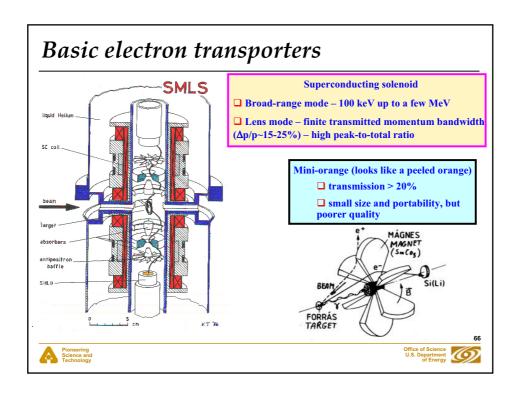


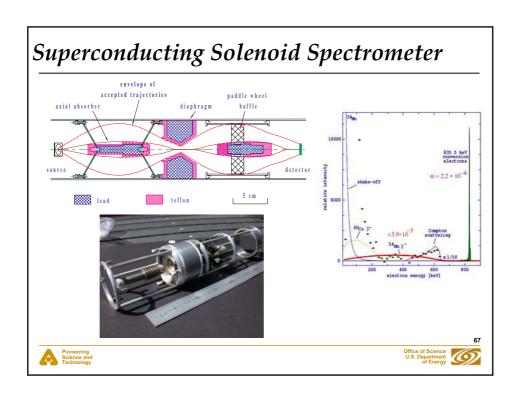


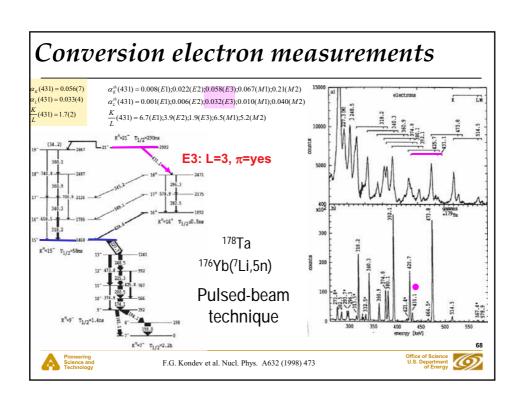


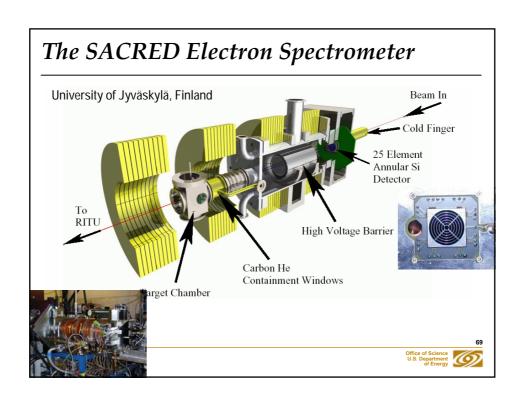


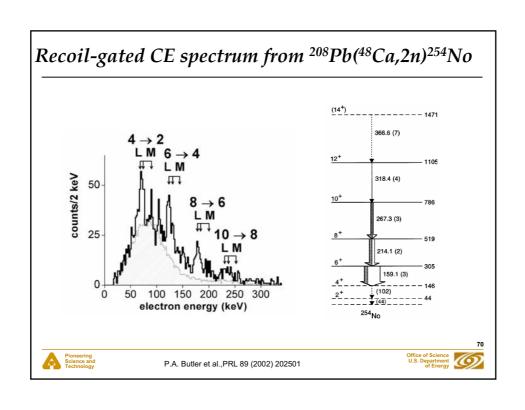


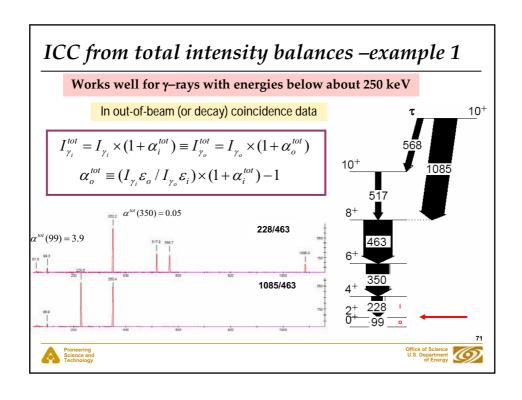


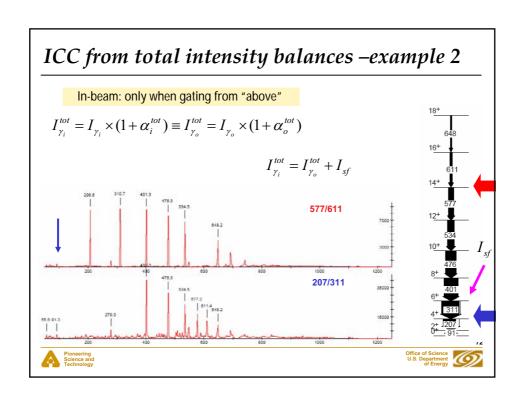










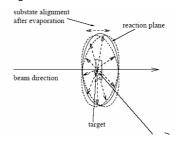


## Angular Distributions

The gamma-rays emitted from nuclear reactions exhibit angular distributions:

$$W(\theta) = 1 + A_{22}P_2(\cos \theta) + A_{44}P_4(\cos \theta)$$
$$A_{22} = \alpha_2 A_2^{\text{max}}; A_{44} = \alpha_4 A_4^{\text{max}}$$
$$\alpha_k(J_i) = \rho_k(J_i) / B_k(J_i)$$

$$\rho_k(J) = \sqrt{(2J+1)} \times \sum_m (-1)^{J-m} < J, m, J-m \mid k0 > P_m(J)$$

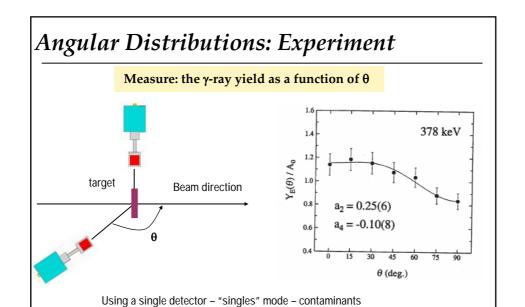


The orientation of the nucleus will be slightly attenuated by the emission of evaporated particles  $(n,p,\alpha)$  and  $\gamma$ -rays.

$$P_{m}(J) = \frac{\exp(-m^{2}/2\sigma^{2})}{\sum_{m=-J}^{J} \exp(-m^{2}/2\sigma^{2})}$$





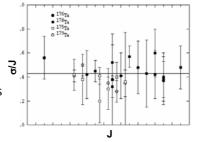


Using a large gamma-ray array – "coincidence" mode - you must be careful!

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## How to determine the mixing ratios?

- l) If both A2 and A4 have been measured see
  E. Der Mateosian and A.W. Sunyar, ADNDT 13 (1974) 407
- II) If only A2 has been measured (A4~0)
  - 1) Determine the attenuation coefficient ( $\alpha_2$ ) for known E2 transitions depopulating levels of known spin (gs band in even-even nuclei)



$$\alpha_2 = A_2^{\rm exp} / A_2^{\rm max}$$

$$A_2^{\rm max} = B_2 \times F_2$$

Tabulated in E. Der Mateosian and A.W. Sunyar, ADNDT 13 (1974) 391

2) For a given transition determine:

$$A_2^{\text{max}} = A_2^{\text{exp}} / \alpha_2$$

3) From E. Der Mateosian and A.W. Sunyar, ADNDT 13 (1974) 407  $get\ \delta$ 







