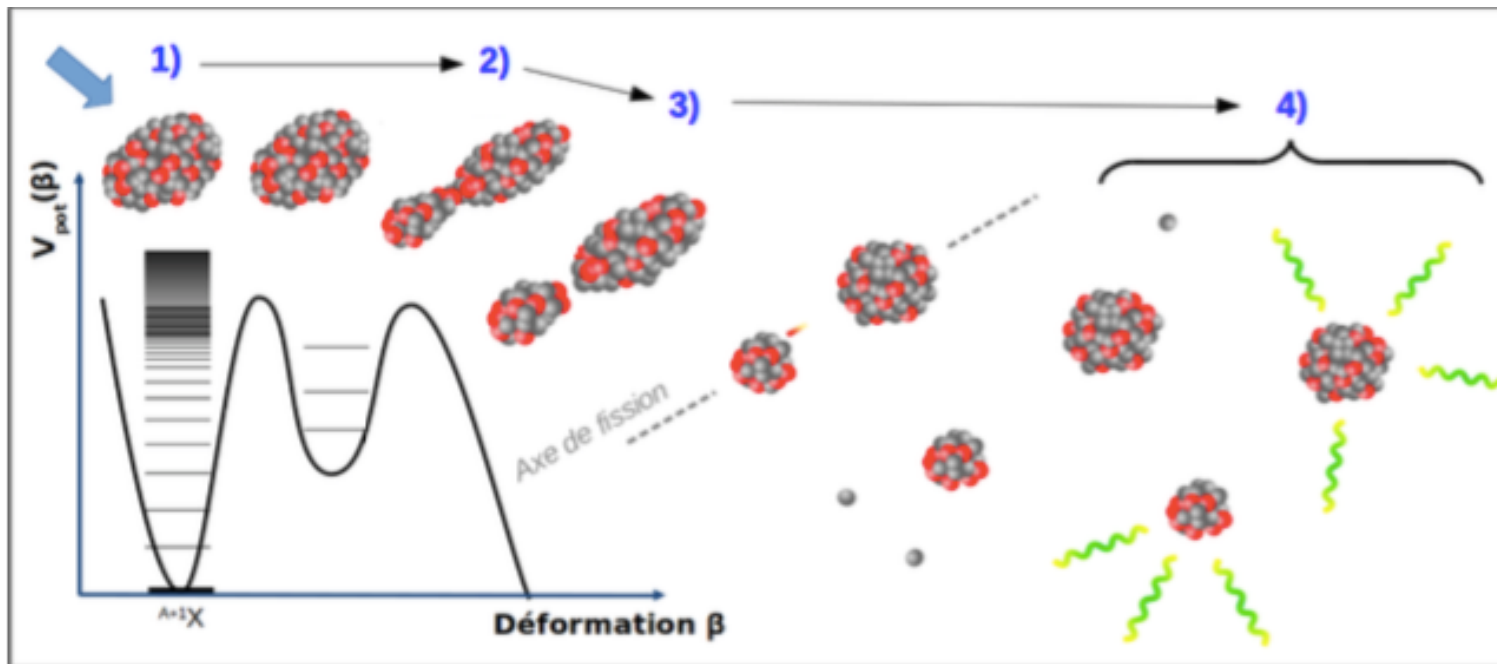


1. Fission process and Fission Yields
  1. Kinetic properties of Fission Fragment
  2. Fission Fragment population of excited states
2. Measurement based on kinematic properties
  1. Energy and time-of-flight based experiments
  2. Rigidity-based experiments
3. Atomic number identification
4. Gamma-spectroscopy techniques
5. Inverse kinematics
  - 5.1 SOFIA experiment
  - 5.2 Multi-nucleon induced fission in inverse kinematics
6. Surrogate reactions
7. Fission yields, systematic uncertainties and errors
8. Conclusions

# 1. Fission Process and Fission Yields

1. Compound Nucleus Formation
2. De-excitation via deformation
3. Saddle point : no-return
4. scission point : formation of fission fragments



Fission **fragments** defined at scission  $A^*$ ,  $Z^*$ ,  $E^*$   
 $E^*$  is released : Fission **product**  $A$ ,  $Z$

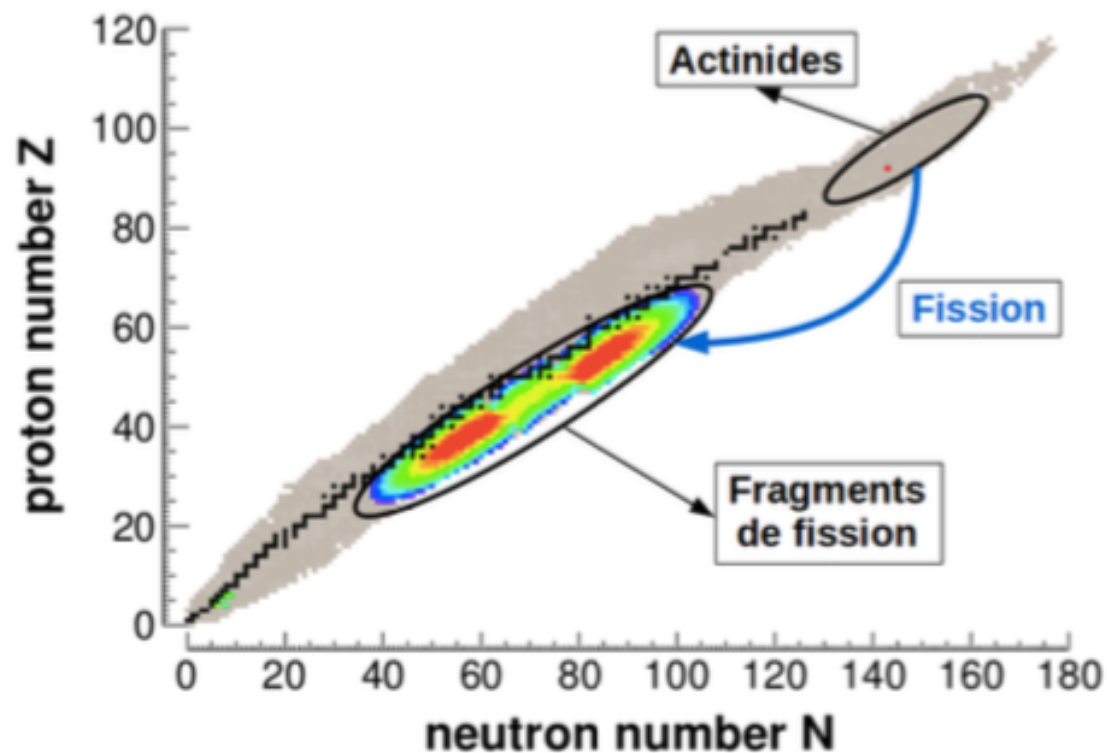
# 1. Fission Process and Fission Yields

Scission point defines the fission fragment properties :

- proton and neutron numbers
- kinetic energy
- excitation energy, angular momentum  
—>(neutron evaporation, gamma emission)

# 1. Fission Process and Fission Yields

## Fission Products are radioactive



Independent yields : **before** beta decay

Cumulative yields : **after** beta decay

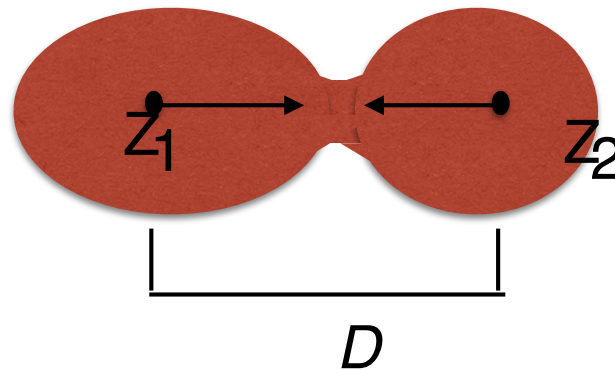


# 1.1 Kinetic properties of fission products

Fission-Fragment kinetic energy defined at scission

$$TKE = 1.44 \frac{Z_1 Z_2}{D}$$

$$D = r_0(A_1^{*1/3}(1 + \frac{2}{3}\beta_1) + A_2^{*1/3}(1 + \frac{2}{3}\beta_2)) + d,$$



(1)

$$TKE = 1.44 \frac{Z_1 Z_2}{D}$$

(2)

$$E_1 = \frac{A_2}{A_1} E_2$$

(3)

$$E_1 + E_2 = TKE$$

# 1.1 Kinematic properties of fission products

**Assumptions :**

**UCD**

**D ~constant**

**no n evaporation**

$$\frac{Z_1}{A_1} = \frac{Z_2}{A_2} = \frac{Z_f}{A_f} = \alpha_f \quad (4)$$

(2;3)

$$E_2 \frac{A_1 + A_2}{A_1} = TKE$$

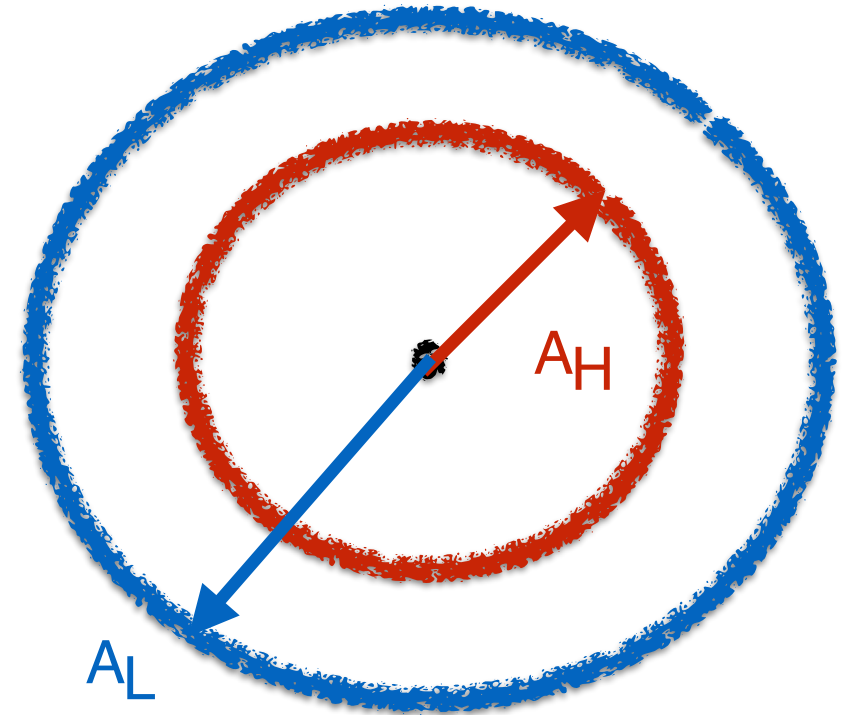
(3;4)

$$E_2 \frac{Z_1 + Z_2}{Z_1} = TKE$$

# 1.1 Kinematic properties of fission products

$$E_2 = \frac{1.44}{D} \frac{Z_1}{Z_1 + Z_2} Z_1 Z_2$$

$$E_1 = \frac{1.44}{D} \frac{Z_2}{Z_1 + Z_2} Z_1 Z_2$$

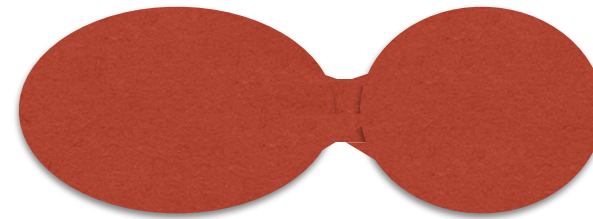


**Light** fragment is **faster**

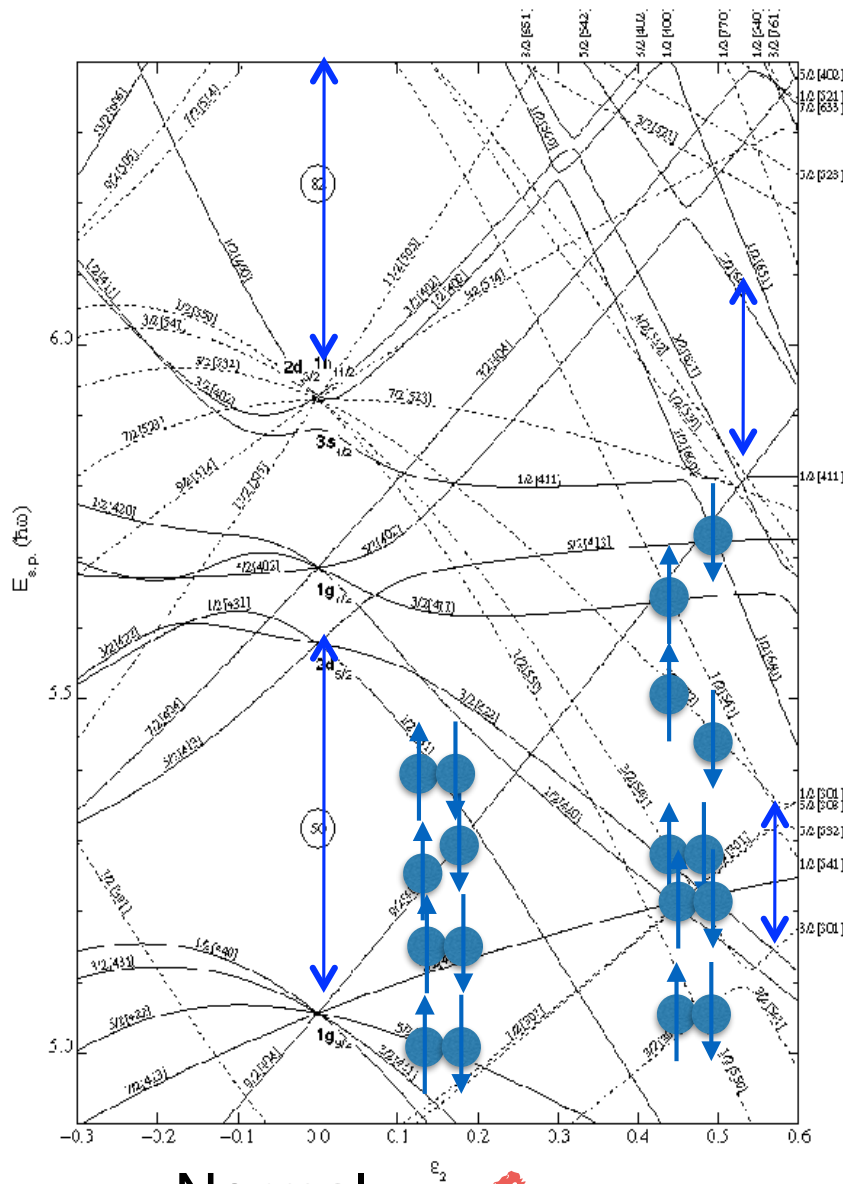
Fission fragments are slow (typically 1 cm/ns  $\Leftrightarrow$  1 A MeV)

They are emitted in  $4\pi$

## 1.2 fission products : population of excited states

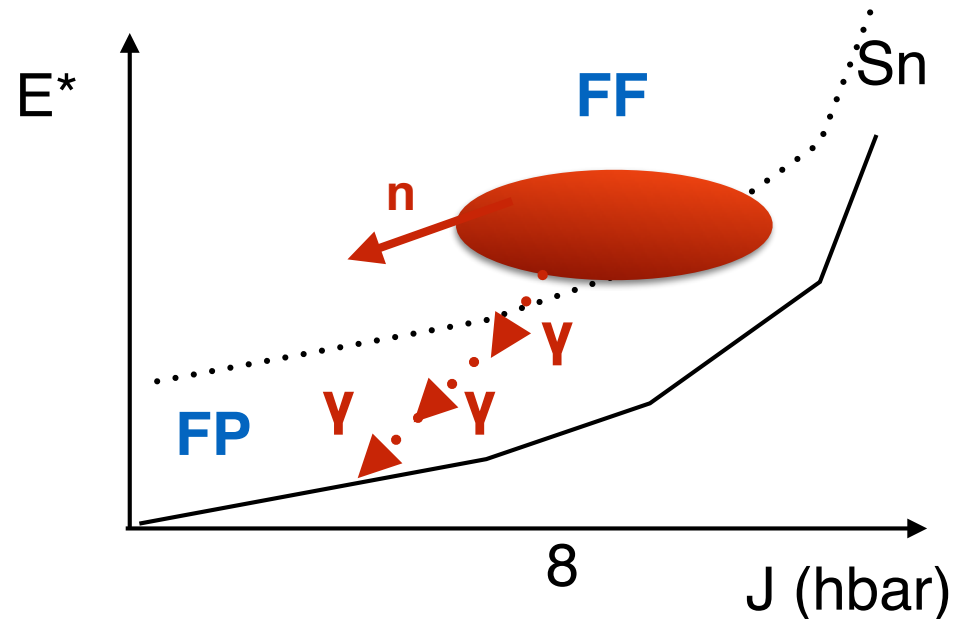


Deformed configuration of fragments  
at scission:  
particle-hole excitation in the level states  
Angular momentum and  $E^*$



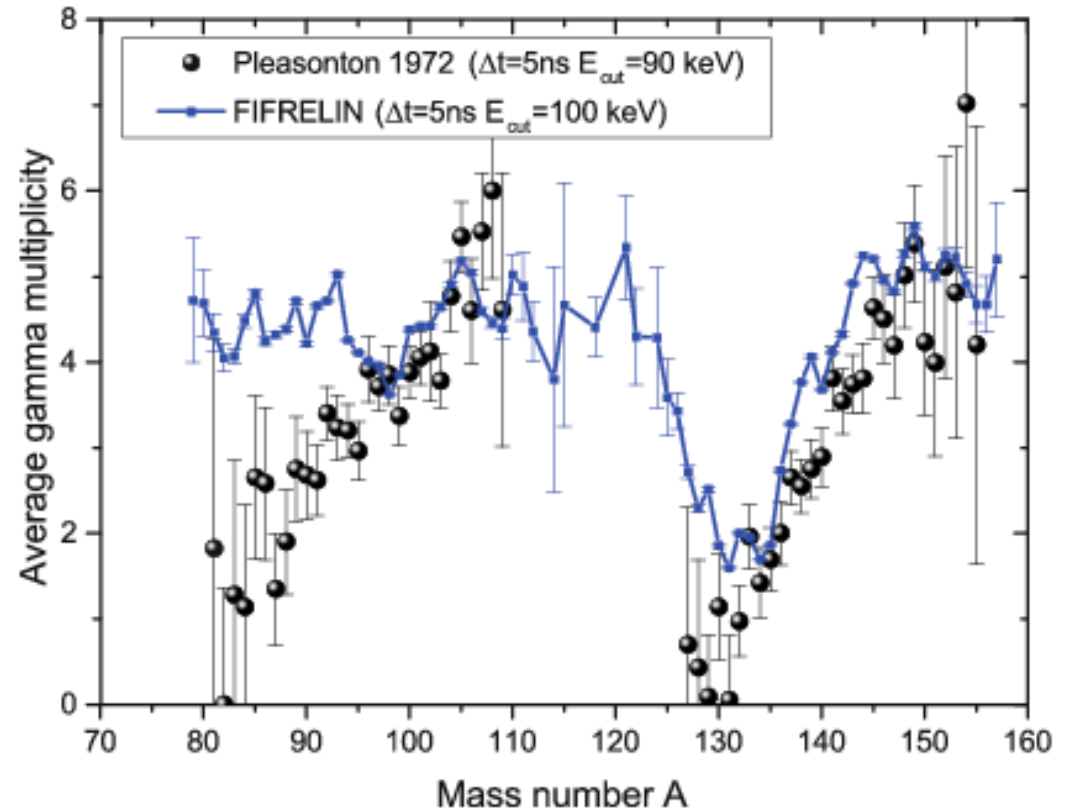
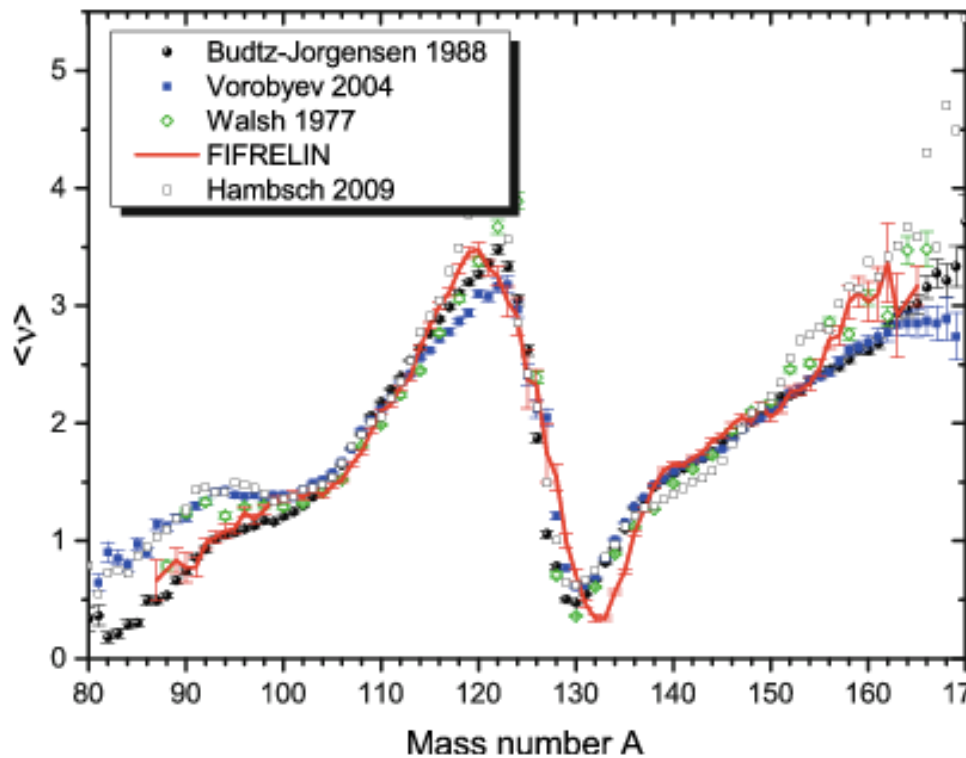
Normal  
deformation

Scission



## 1.2 fission products : population of excited states

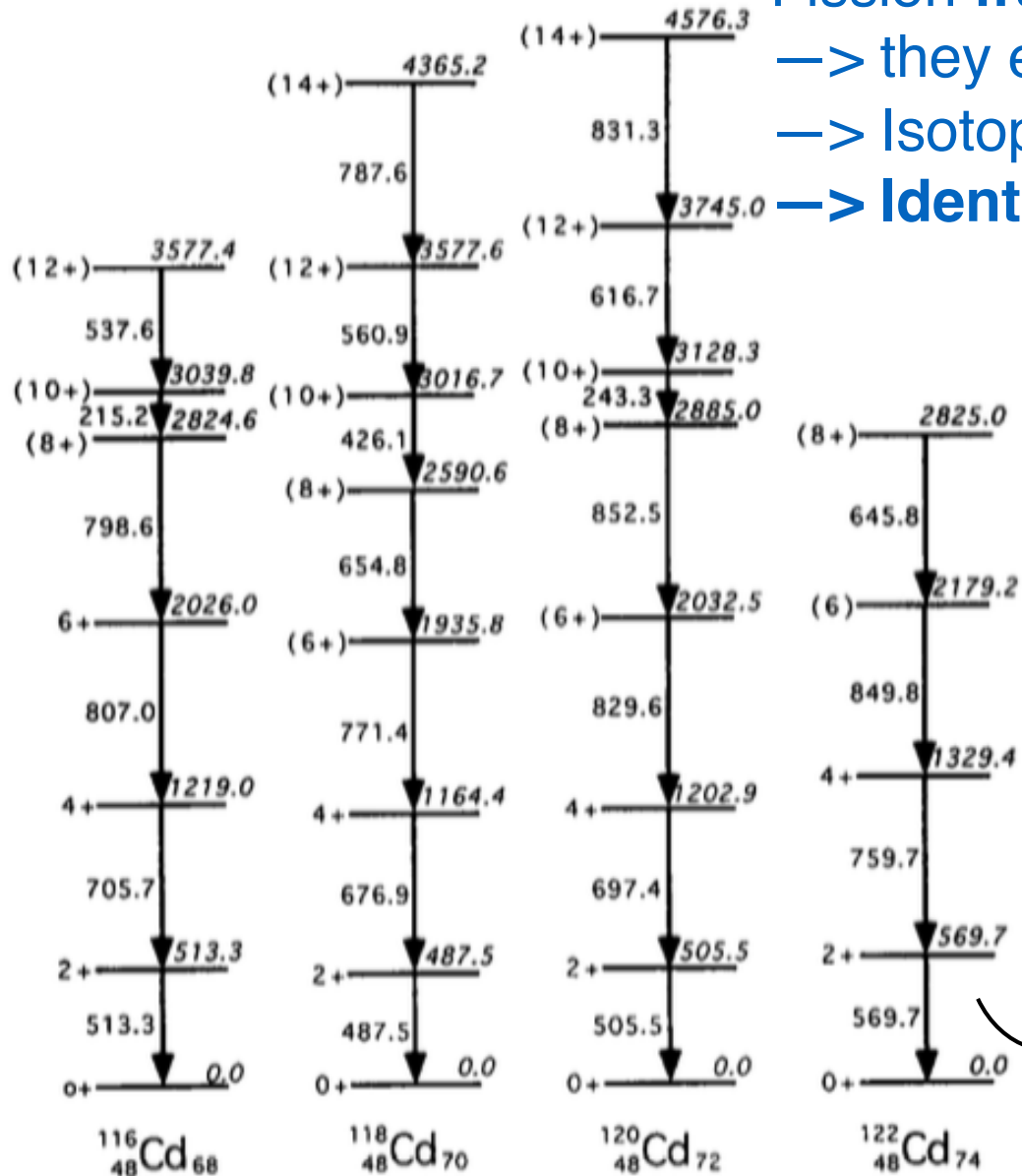
$\nu(A)$  and  $\gamma(A)$  :: excitation energy is depending on the sorting



From O. Litaize et al., EPJA 51(2015)

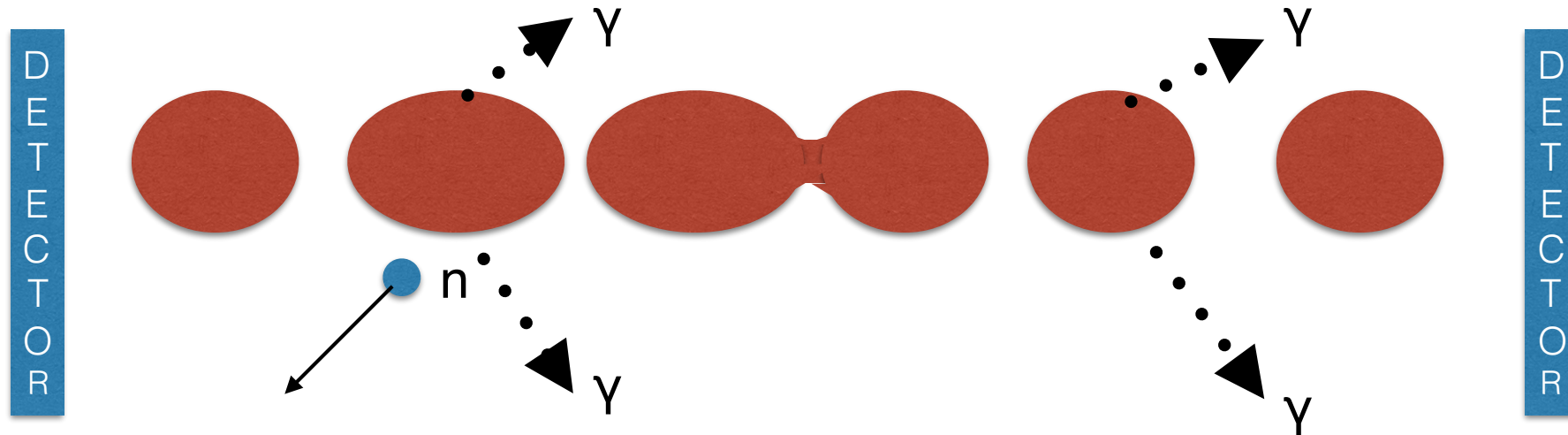
## 1.2 De-excitation properties of fission fragments

J. H. Hamilton *et al.*



Fission **fragments** are produced in excited states  
 —> they emit gamma rays  
 —> Isotopic identification of fission **products**  
 —> **Identification of the fission products pair**

## 2. Measurements based on kinematical properties



Fragment mass identification

$$A_1^* \beta_1^* c = A_2^* \beta_2^* c$$

$$A_1^* + A_2^* = A_f$$

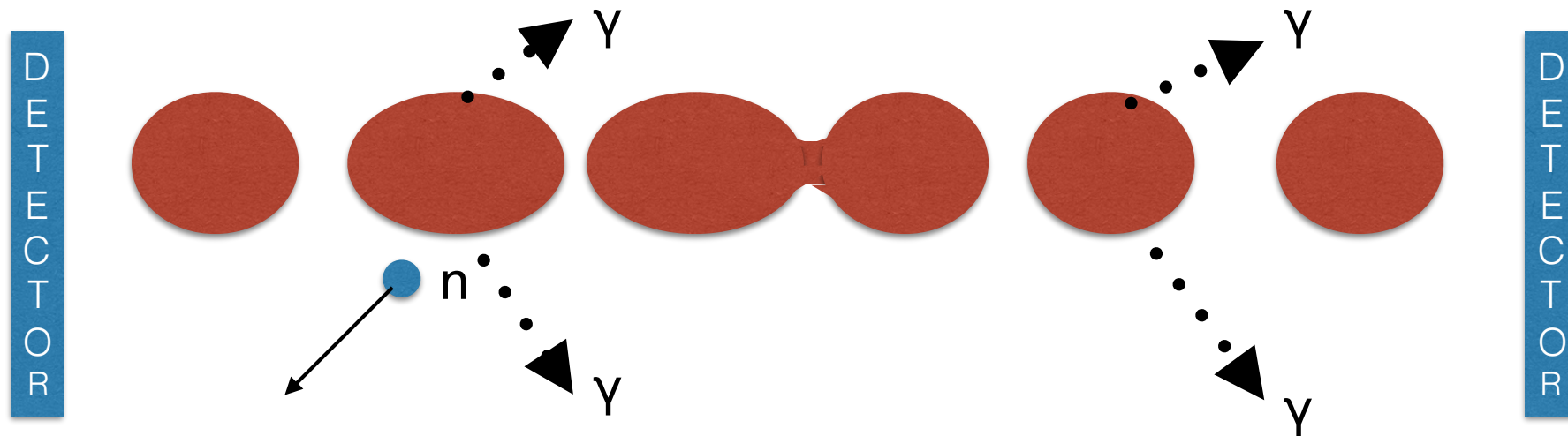
Momentum and Mass conservation

$$\langle \beta_{1,2}^* \rangle = \langle \beta_{1,2} \rangle$$

Isotropic emission of neutrons

$$A_1^* = A_f \frac{\beta_2}{\beta_1 + \beta_2}$$

## 2. Measurements based on kinematical properties



**Product** mass identification :

$$E_{1,2} = \frac{1}{2} A_{1,2} m_0 \beta_{1,2} c^2$$

Energy and velocity measurement

$$A_{1,2} = \frac{2E_{1,2}}{2A_{1,2} m_0 \beta_{1,2} c^2}$$

$$v_{1,2} = A_{1,2} - A_{1,2}^*$$

neutron multiplicity



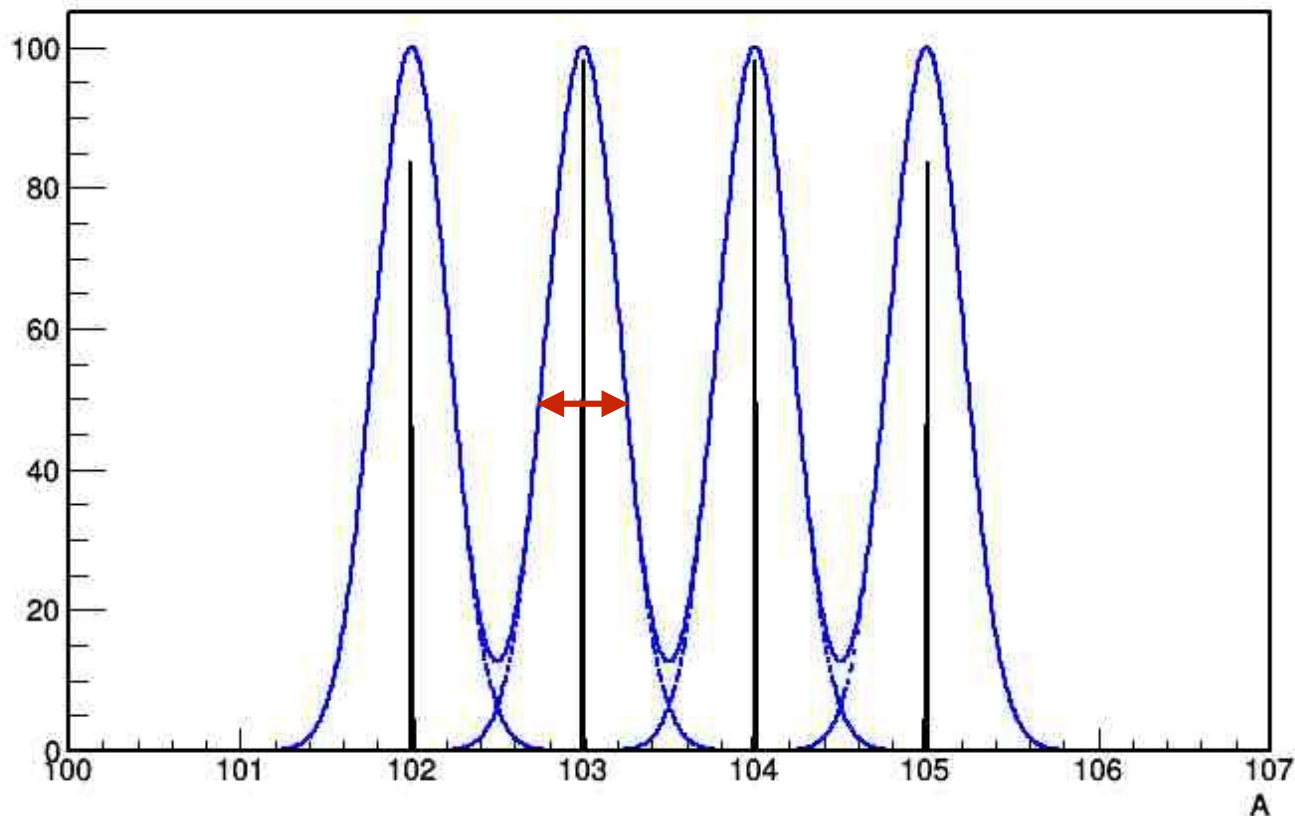
## which resolution ?

Resolution of 1uma does depends on the mass range :

Around  $A \sim 100$  : 2 adjacent masses are distant by 1%

Around  $A \sim 50$  : 2 adjacent masses are distant by 2 %

Around  $A \sim 150$  : 2 adjacent masses are distant by 0.7%



FWHM = 0.5 %

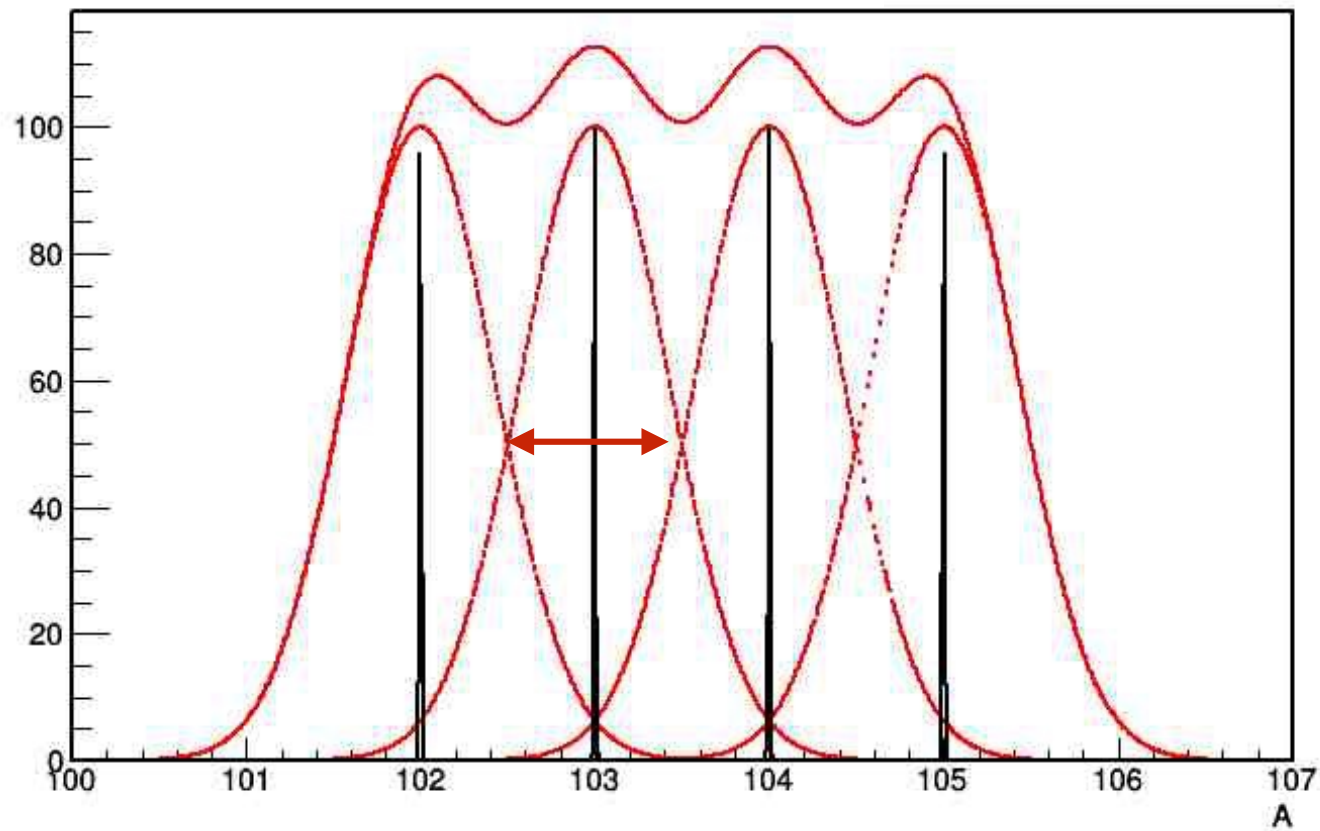
Resolution is important for a better counting

## which resolution ?

$DA/A \sim 1 :: \text{Resolve } A \sim 100$

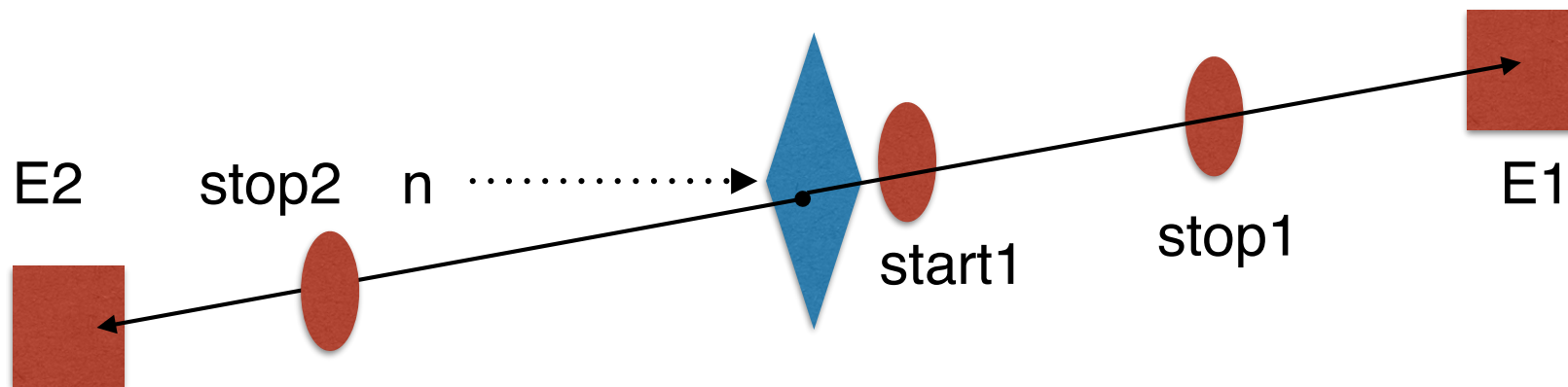
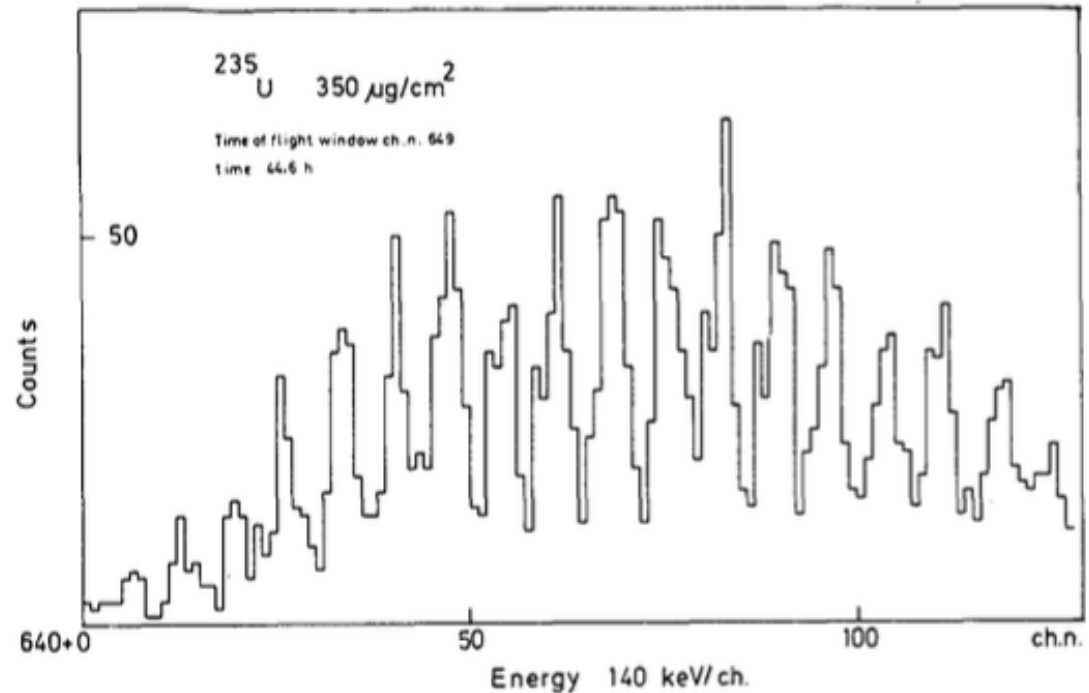
$DA/A \sim 0,5 :: \text{Resolve } A \sim 200$

$\text{FWHM} = 1 \%$



## 2.1 Energy and Time-of-Flight based experiments

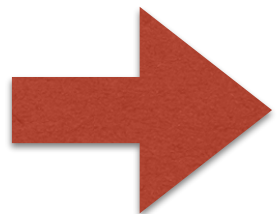
$$\frac{\partial A}{A} = \sqrt{\frac{\partial E^2}{E} + 2 \frac{\partial \beta^2}{\beta}}$$



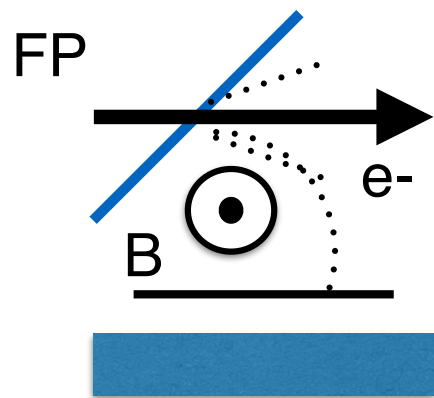
BEST : 0,6% in COSIFANTUTE, ILL, 1980

# Time-of-Flight measurement

Minimum layer  
Fast timing



Secondary electron detectors



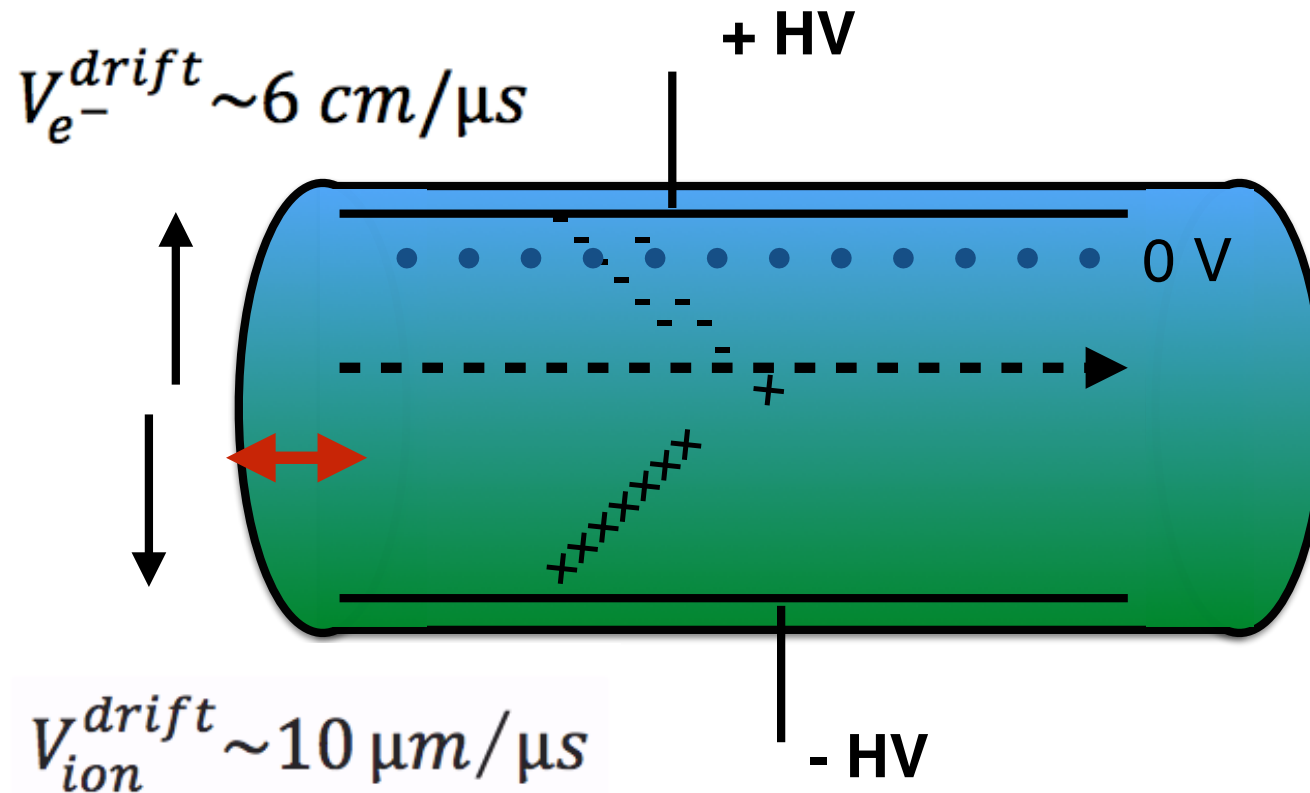
+HV  
-HV

MICRO-CHANNEL PLATE

MULTI-WIRE PROPORTIONAL COUNTER

Typical resolution 150ps to 200ps; Typical flight path 50ns  $dT/T \sim 4\%$

# Energy measurement in Ionisation Chambers



$$N_e - W_e = \Delta E$$

$$W_e \sim 15 \text{ eV}$$

$$N_{e^-} \sim 10^7$$

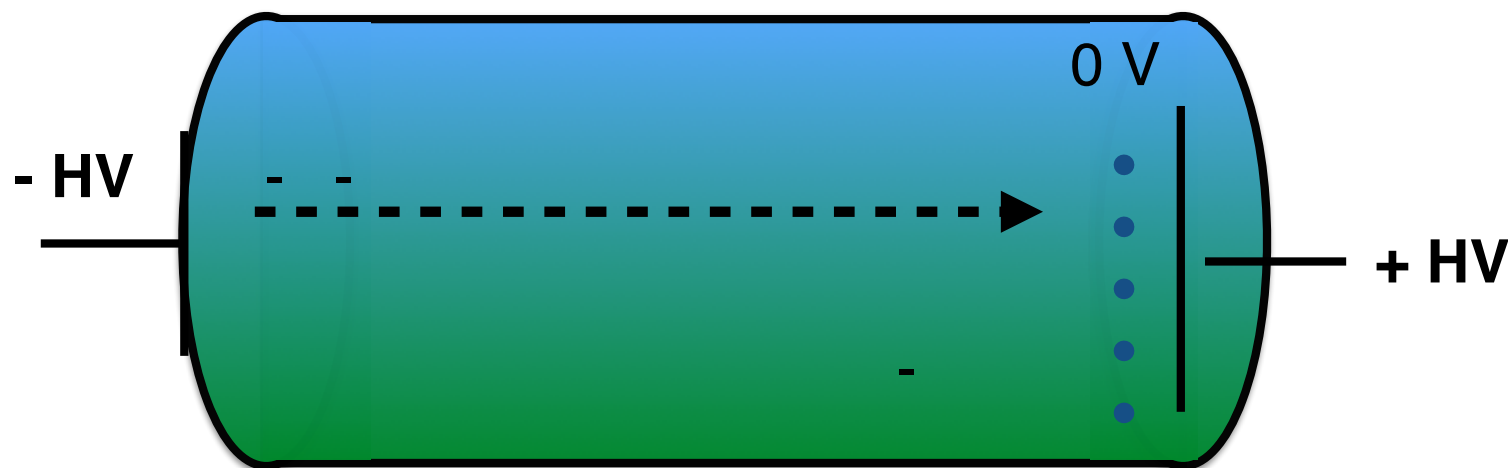
Number of electrons is large  $\rightarrow$  good statistical resolution

Despite drift velocity of ions is slow, they may disturb the apparent total charge collected on the anode  $\rightarrow$  Frisch Grid (0V) to isolate the anode from the positive current



Zone of electric field distortion and energy-loss uncertainty that decreases significantly the resolution

# Energy measurement in Ionisation Chambers



In axial chamber, all electrons are collected on the anode  
Energy-loss collection with 1% precision  
Electronics (noise and amplification) limits to the percentage limit

# Fission-fragment distributions : experimental challenges

Improvement of technology :

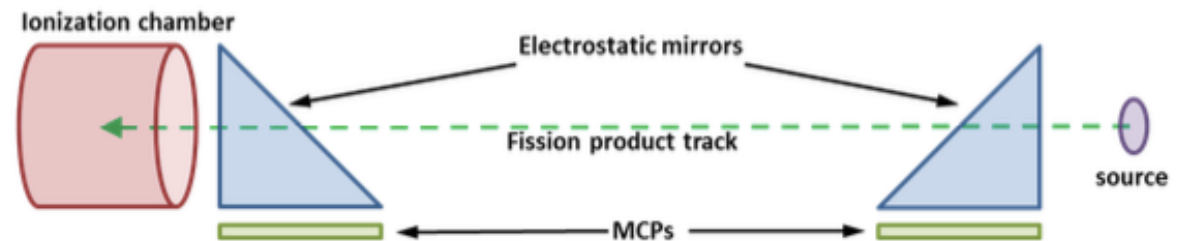
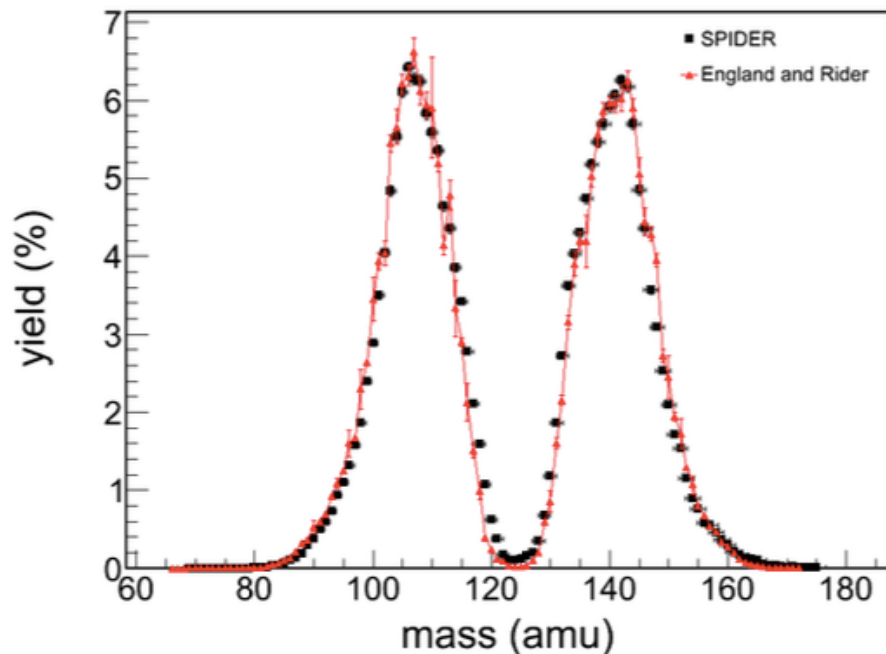
thin layers

SED : amplification in gaz : large surfaces

Digital electronics : energy-loss profile

Physical limits to the resolution :

Energy-loss straggling in materials (stochastic process)

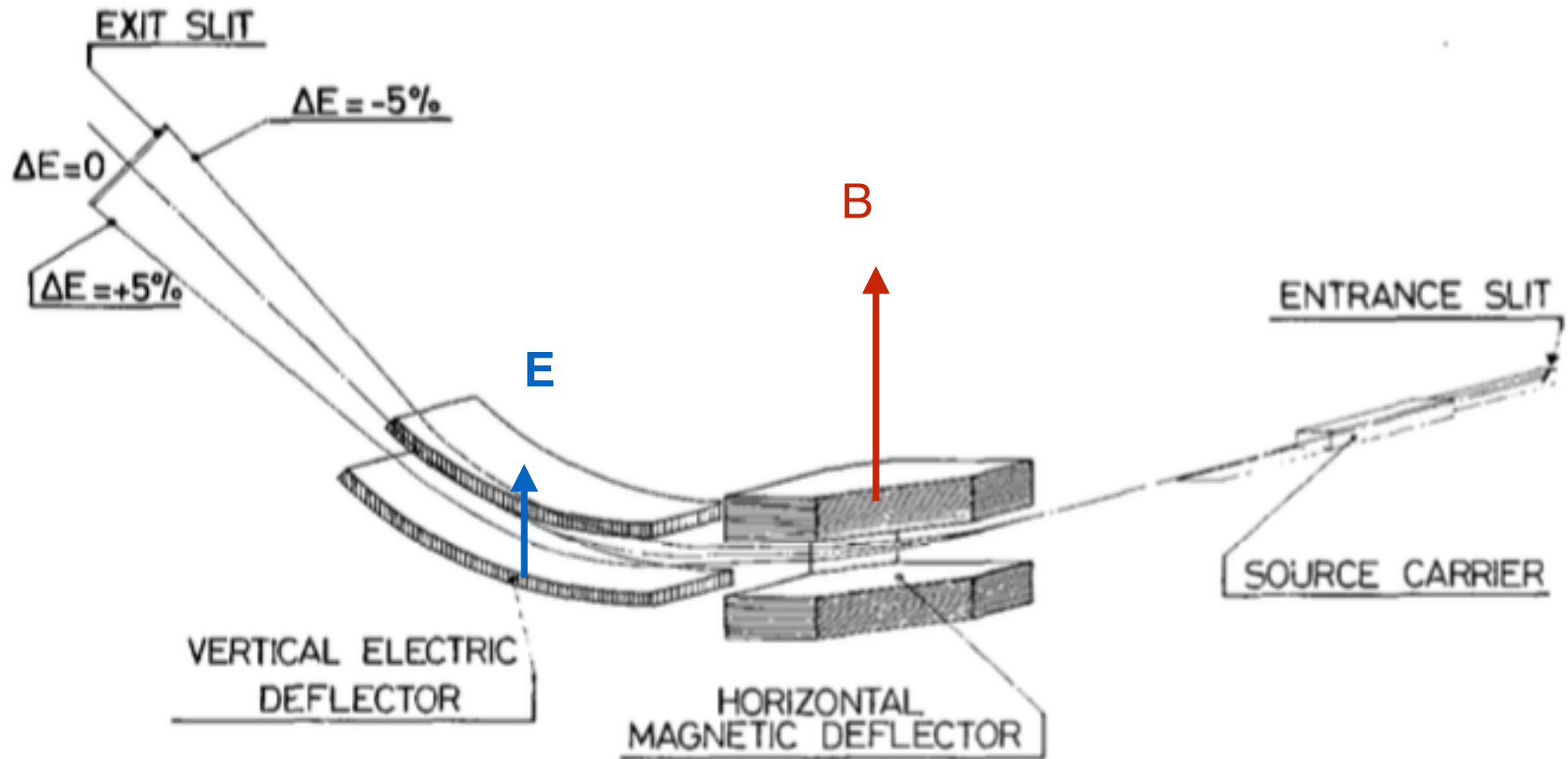


K. C. Meierbachtol, F.K.E Tovesson et al. LA-UR-15-20101

# Rigidity-based experiment

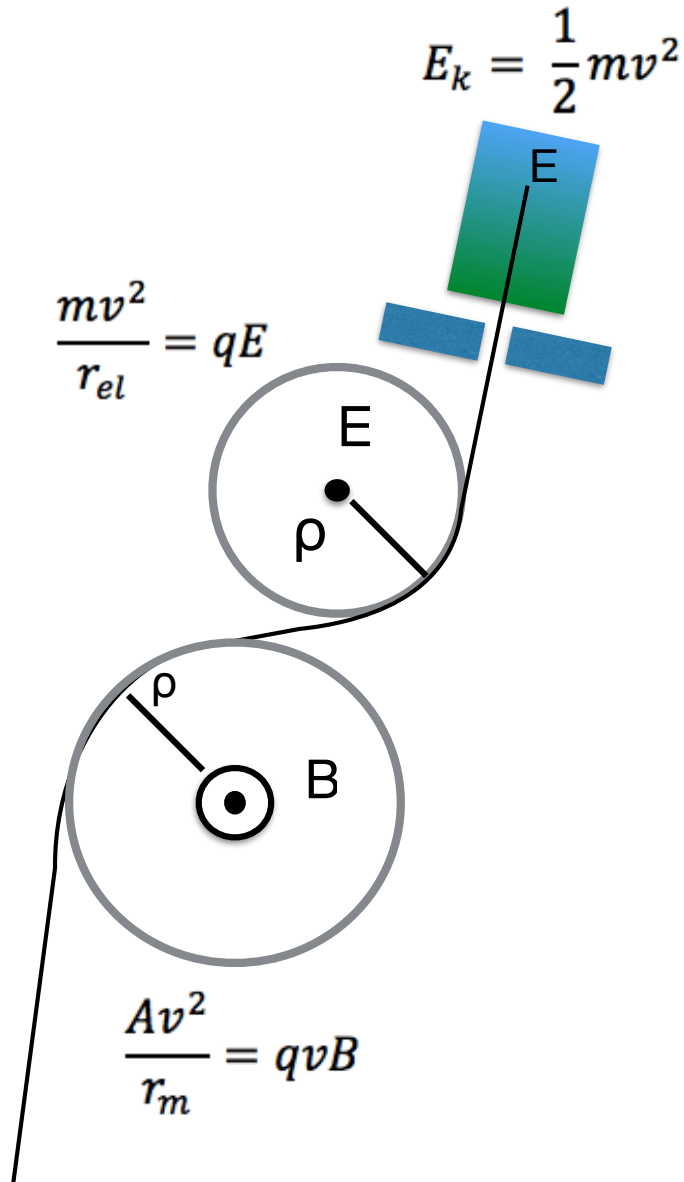
## LOHENGRIN spectrometer

No ToF measurement !





# Rigidity measurement : Lohengrin

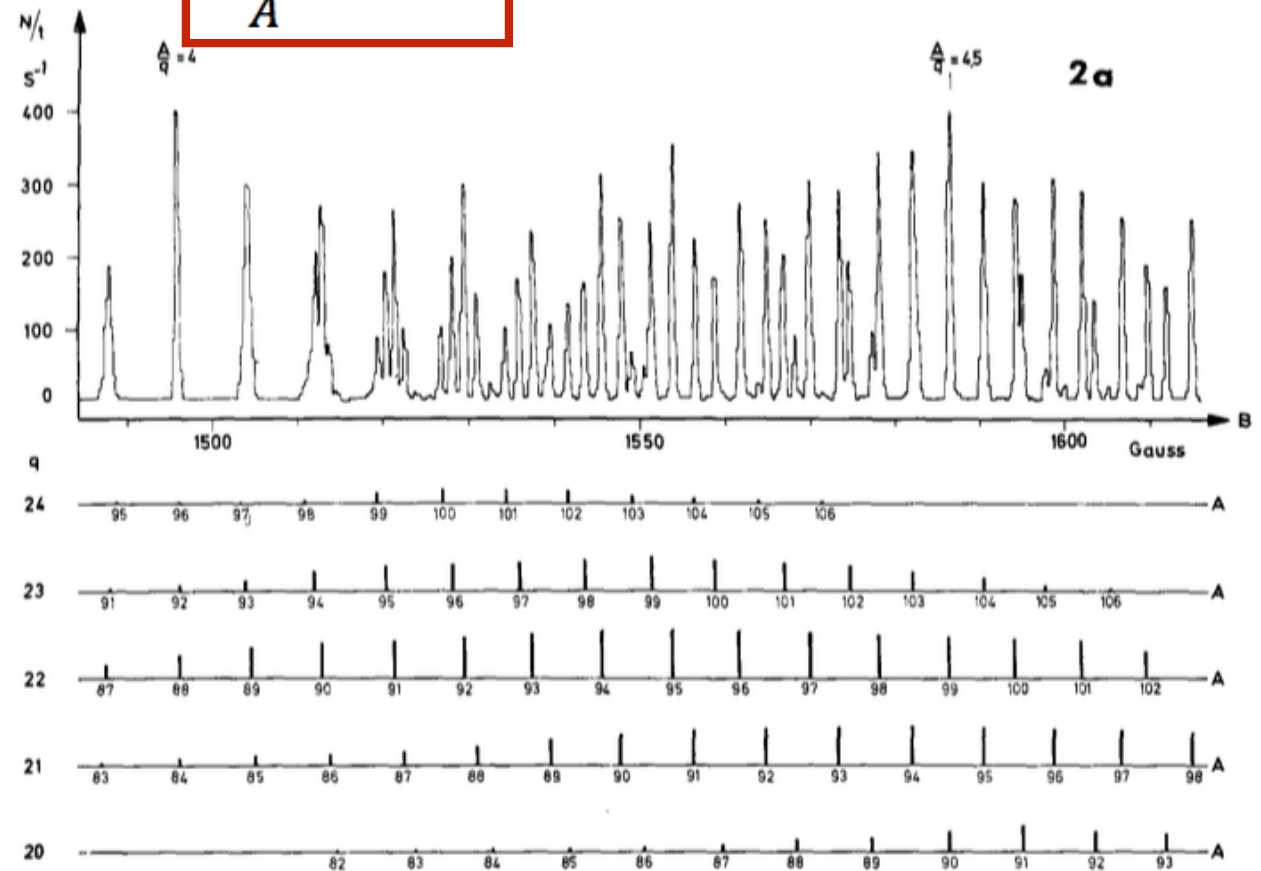


$$\frac{A}{q} \frac{Av^2}{q} = r_m^2 B^2$$

$$\frac{A}{q} = \frac{r_m^2 B^2}{r_{el} E}$$

$$\frac{E_k}{q} = \frac{1}{2} r_{el} E$$

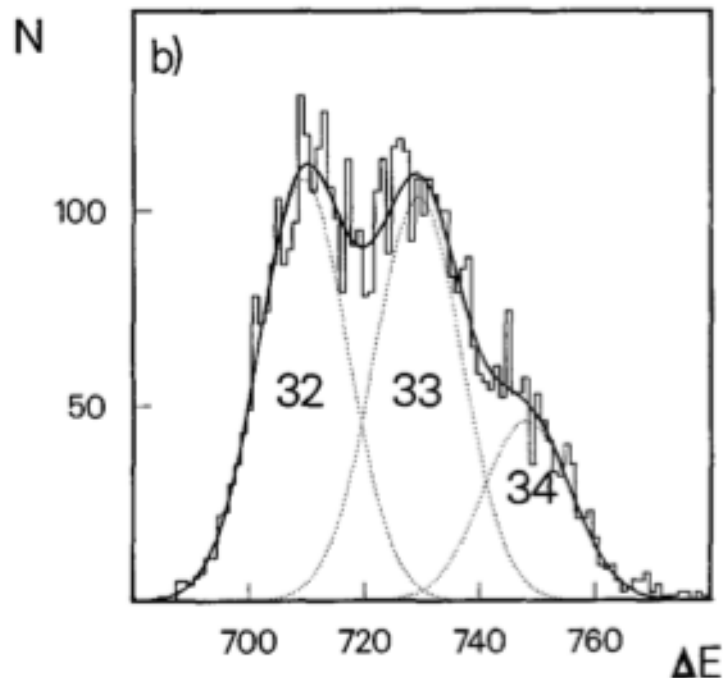
$$\frac{\partial A}{A} \sim 0.01 \%$$



## 3. Atomic Number identification

### Energy loss in ionisation chamber

$$-\frac{dE}{dx} = 2\pi N_a r_e^2 m_e c^2 \rho \frac{Z}{A} \frac{q^2}{\beta^2} \left[ \ln \left( \frac{2m_e \gamma^2 v^2 W_{max}}{I^2} \right) - 2\beta^2 - \delta - 2\frac{C}{Z} \right]$$



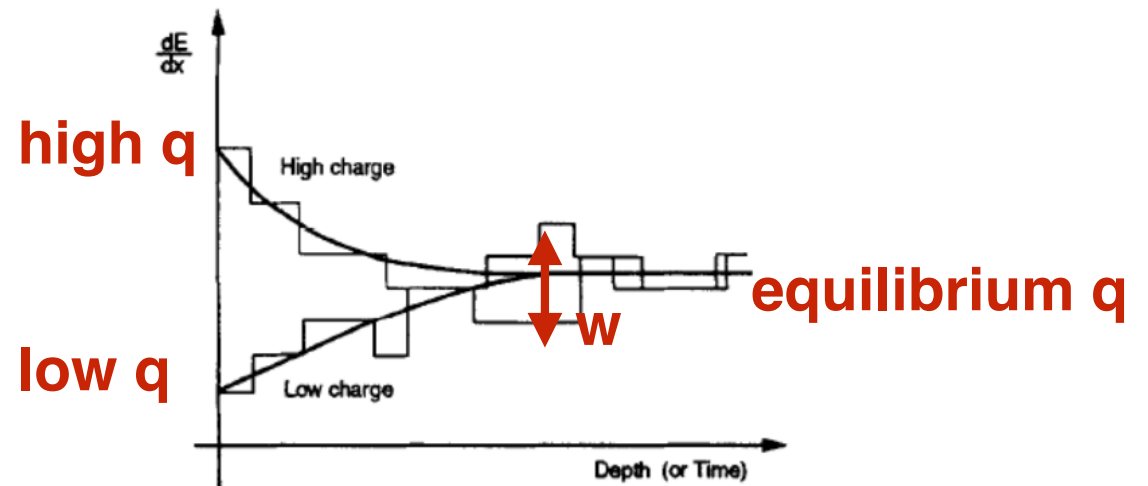
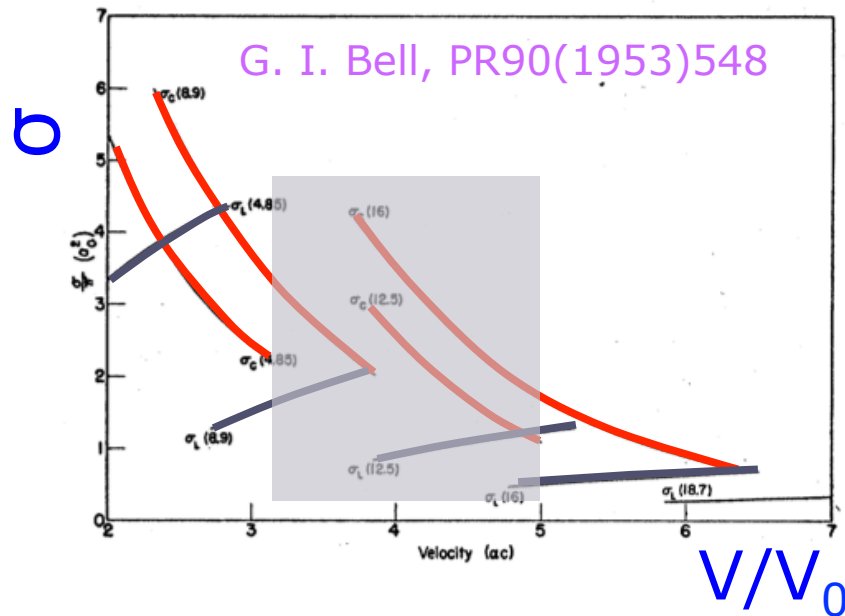
Fission products  
 have a broad q-state distribution  
 —> energy-loss resolution is limited

### 3. Atomic Number identification

#### Energy-loss of ion in material

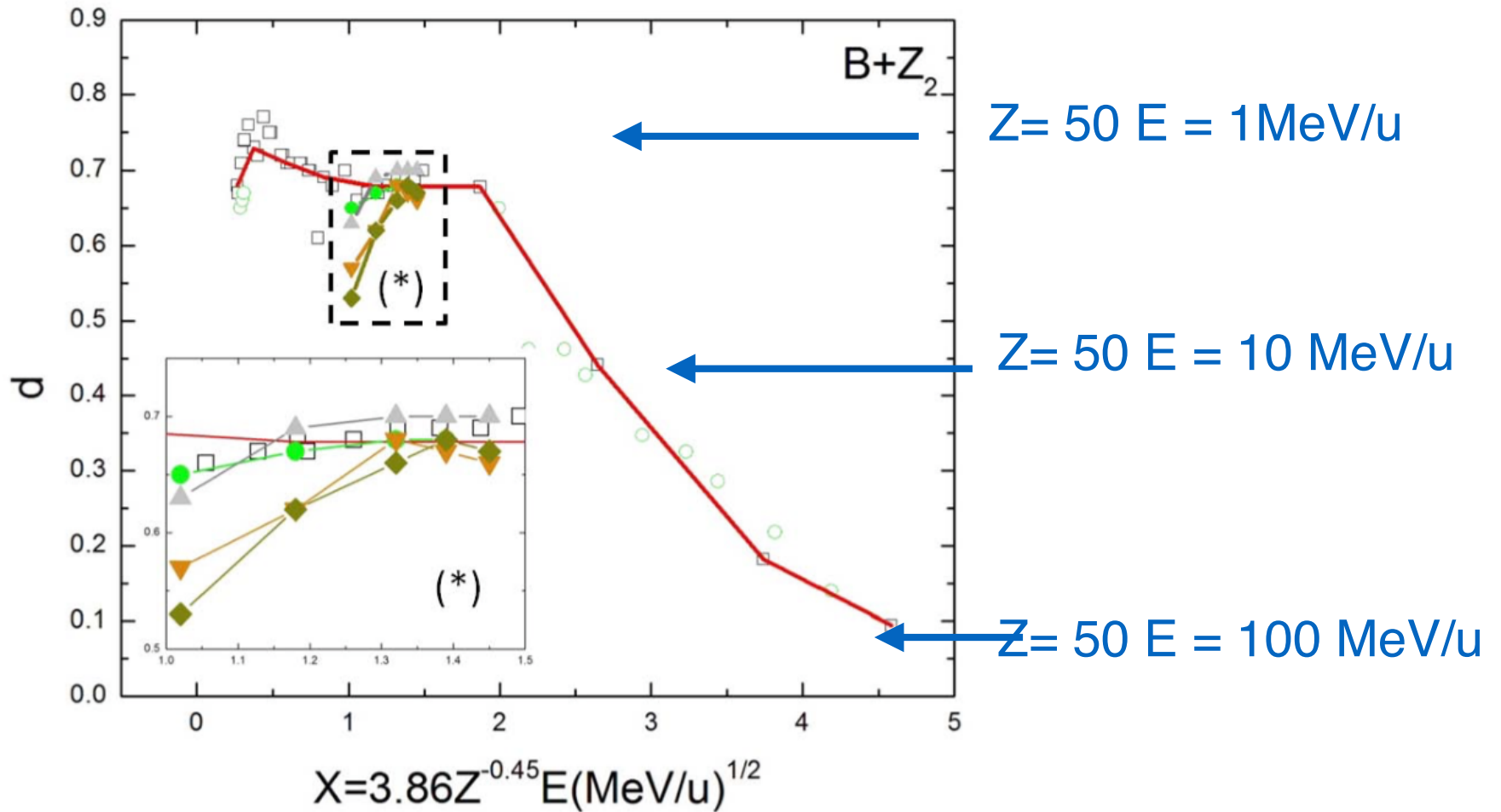
Energy-loss prop. to ionic charge state

Electron loss and capture cross sections depend on ion energy :

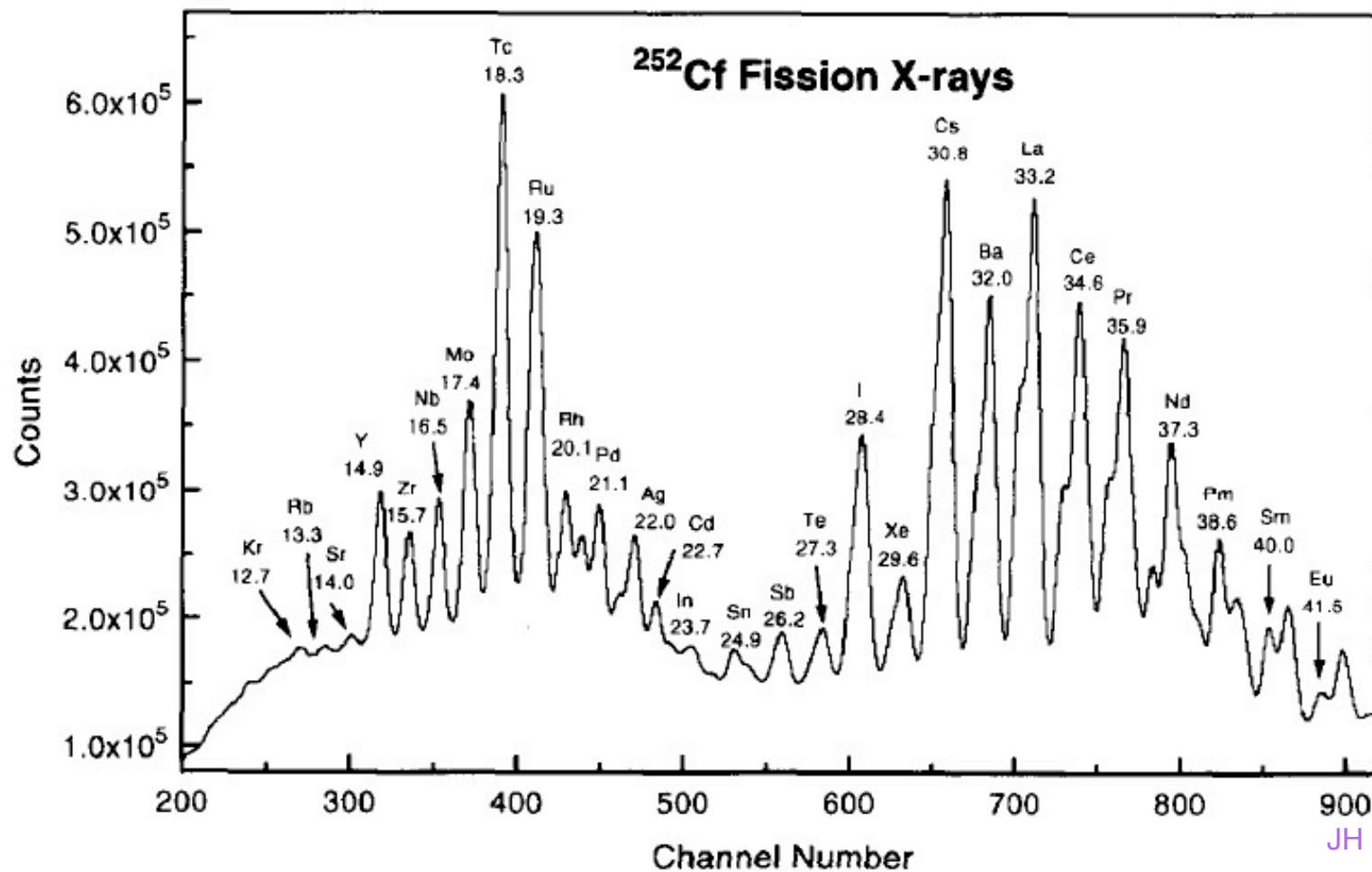


# ionic charge dispersion diminishes with energy

W



## Detection of X-rays



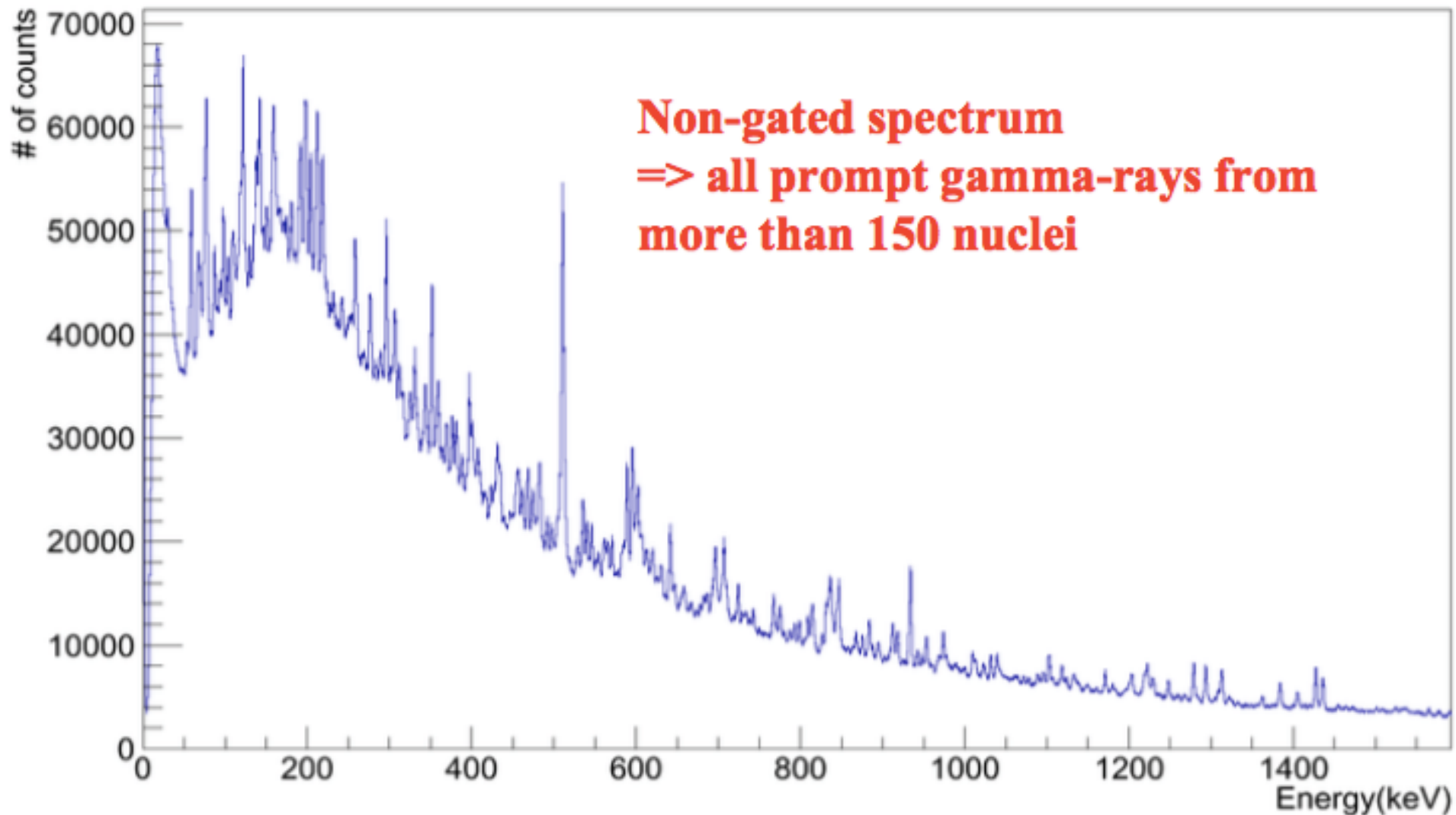
JH Hamilton Prog. Part. Nucl. Phys. 35

small detection efficiency —> Difficulty to connect with mass distribution

## 4. Isotopic identification from gamma spectroscopy

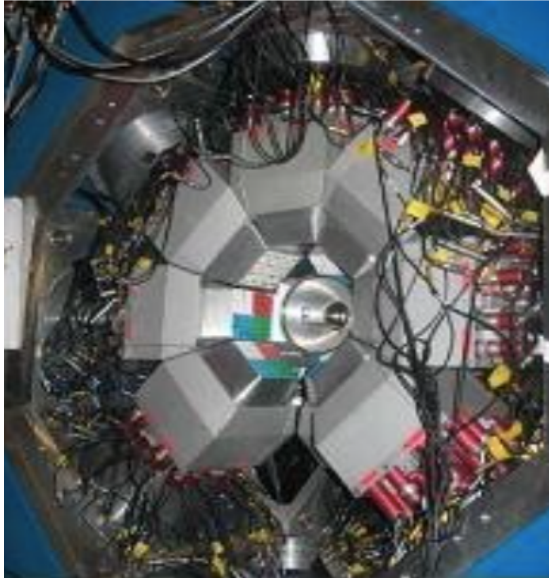
Fission products may be identified unambiguously from their  $\gamma$ -ray emission

Difficulties : Large background from beta decay

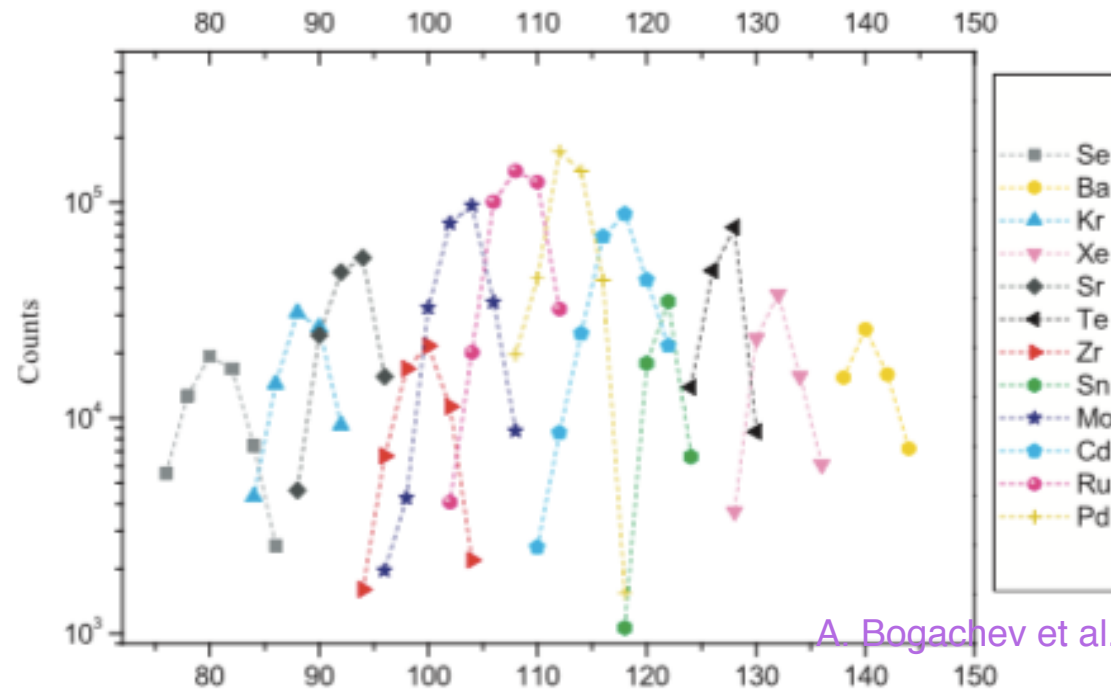
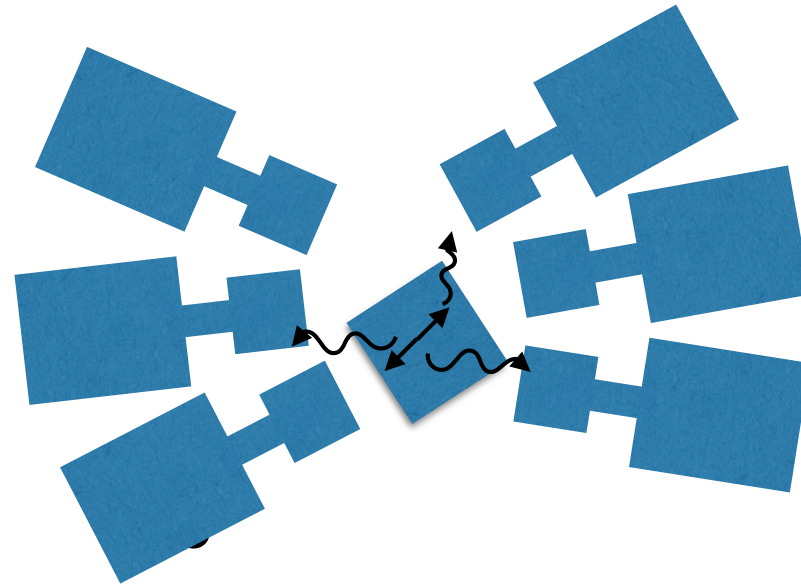


# Isotopic identification from gamma spectroscopy

$\gamma$ - $\gamma$  technique :



EXILL, ILL



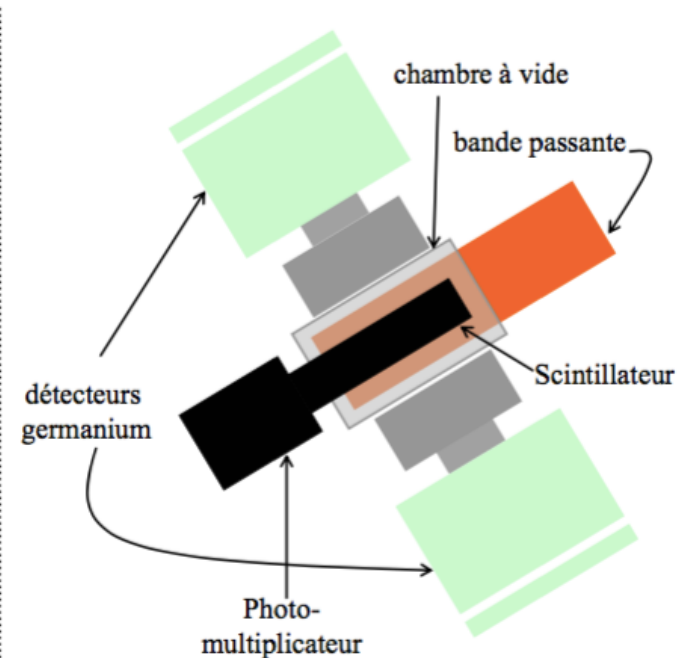
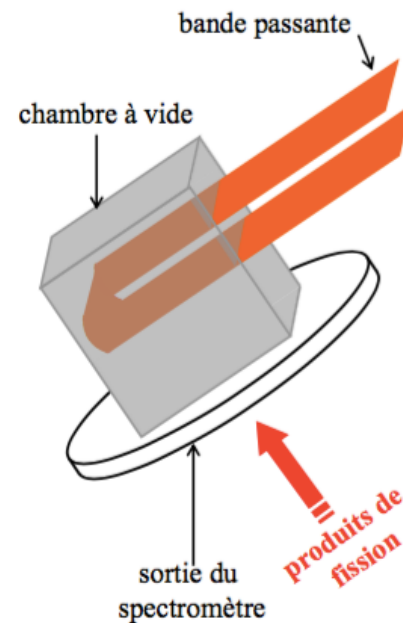
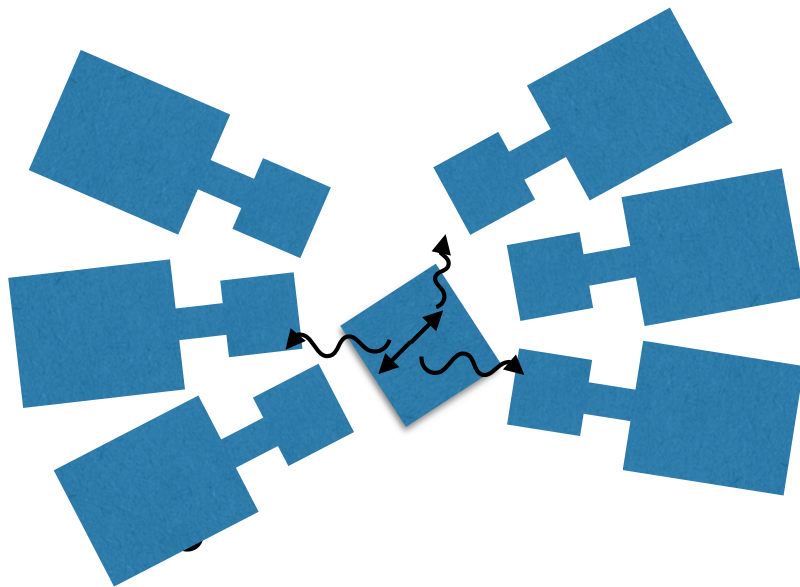


## 4. Isotopic identification from gamma spectroscopy

To fight against background :

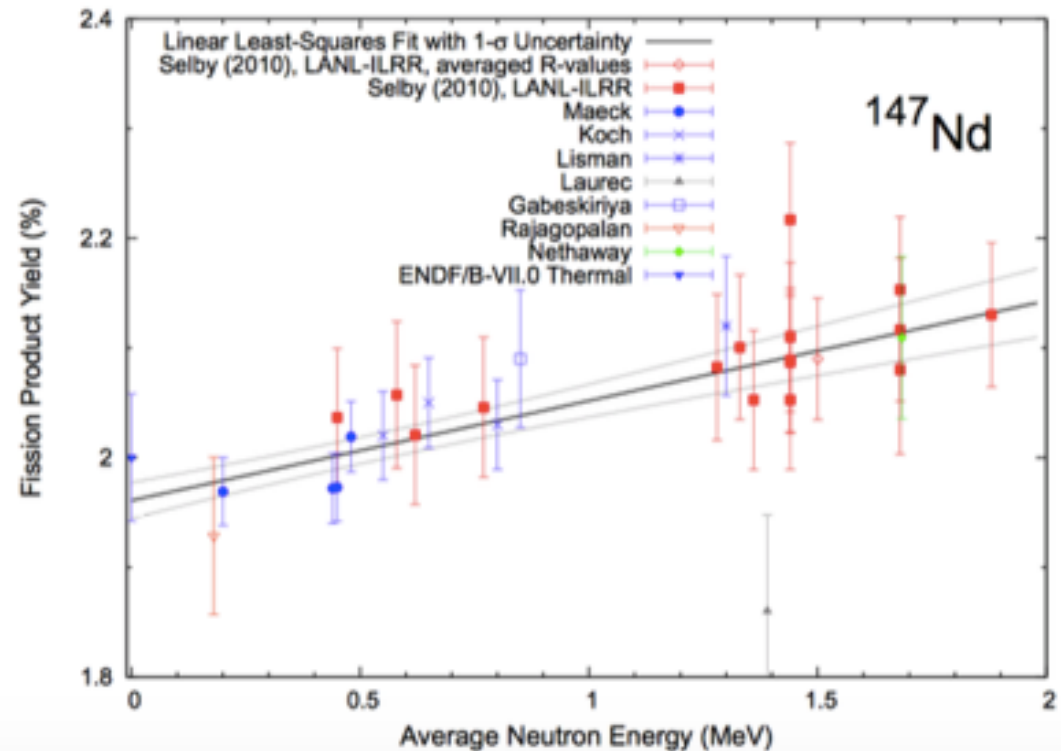
- larger number of detectors ( $\gamma\gamma\gamma$  technique)
- external trigger experiment  
(fast ionisation chamber  $\rightarrow$  thin target)

–  $\beta\gamma$  coincidence experiment  
( access to cumulative yields)

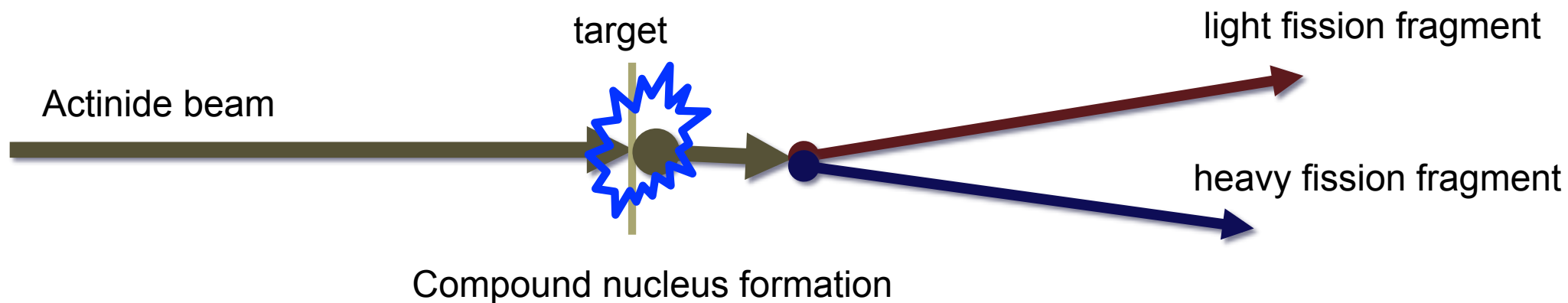




==> Difficult measures !!



# Inverse kinematics : need for accelerator complexe

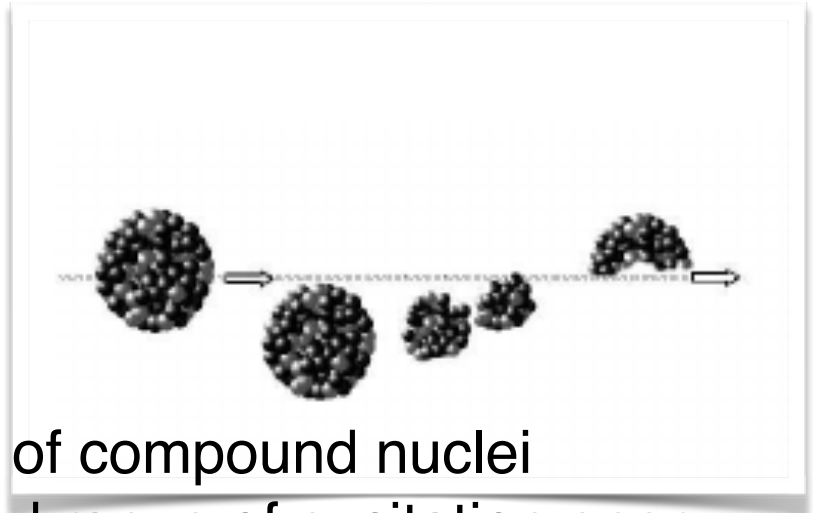
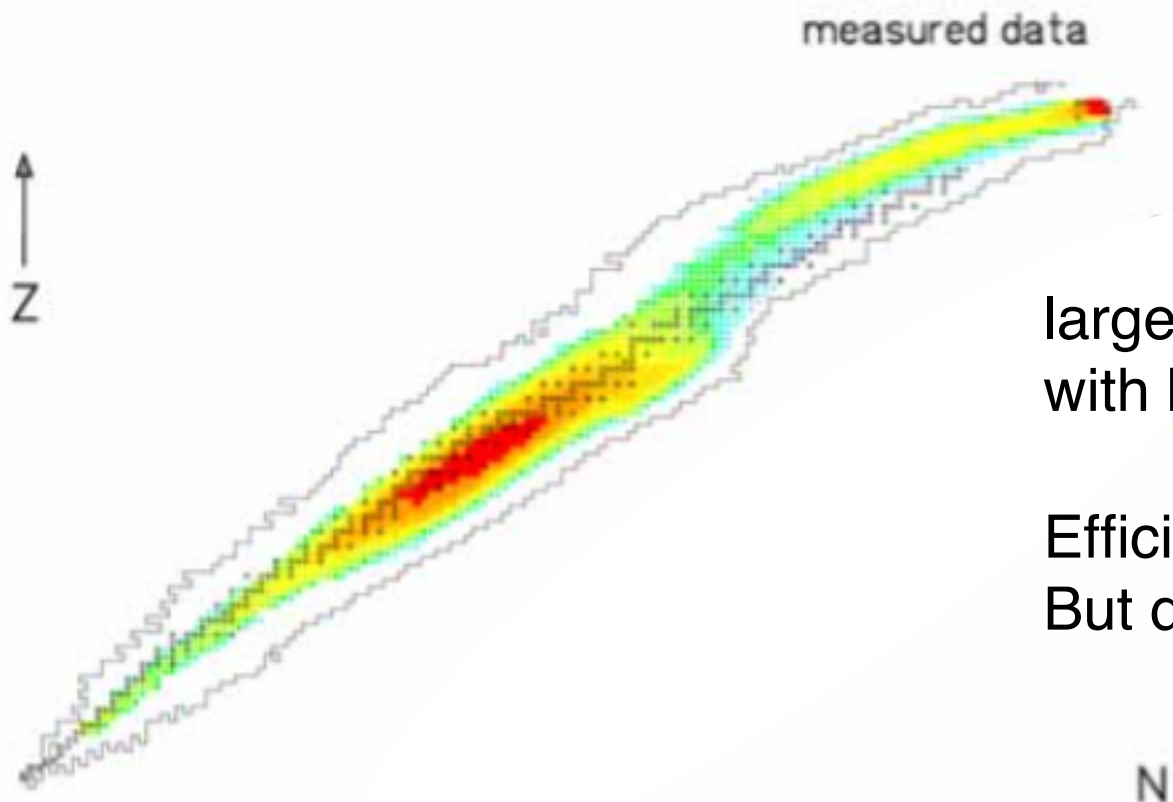


## Increasing fission-fragment velocity for a better resolution

- + forward focusing for a better detector acceptance
- time-of-flight resolution decreased

Different regimes : relativistic energy  
Coulomb barrier energy

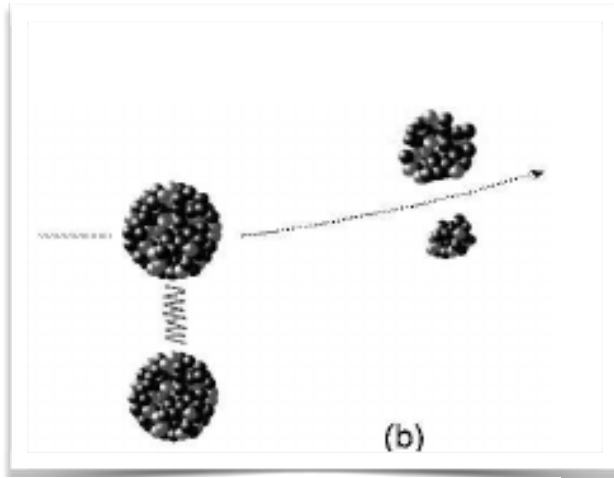
Relativistic energy : fragmentation or spallation



large set of compound nuclei  
with broad range of excitation energy

Efficient to produce RIB  
But difficult to study fission process !

## Relativistic energy : Coulomb excitation

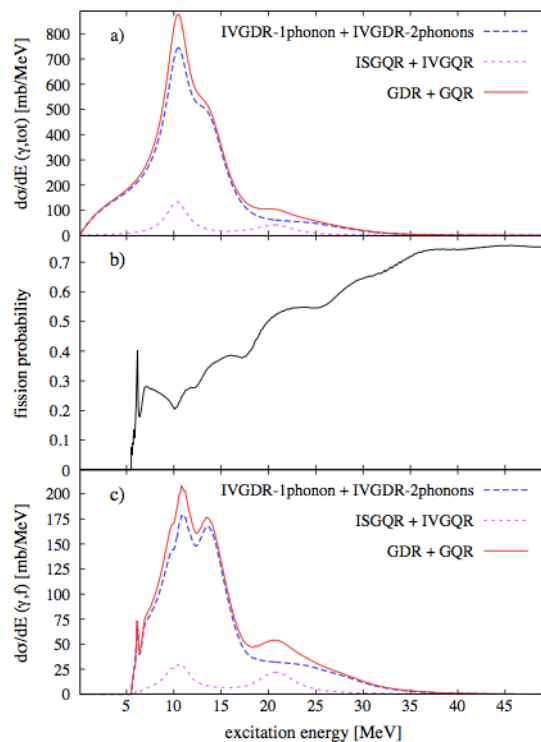


To select Cb excitation events :

- large-Z target
- $Z_1 + Z_2 = Z_{\text{beam}}$
- subtraction of low-Z target induced fission

Relativistic kinetic energy

Moderate excitation energy !



How to produce actinide beams ?

Natural U allows to extract  $^{238}\text{U}$ , but  $^{235}\text{U}$

Th is possible ...

But actinides are very radioactive and it becomes very costly to decontaminate an accelerator !!

Two steps experiments :

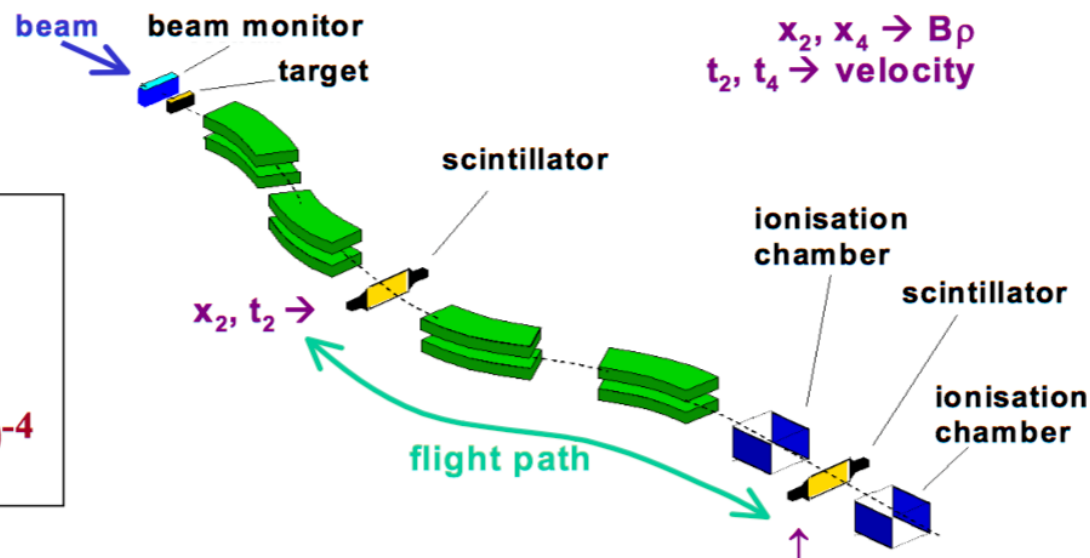
1) production of actinides

2) induce fission and detect fission fragments

# In-flight production of actinides

- 1) Fragmentation of U beam on target
- 2) Use of separator

$$\begin{aligned}
 A / \Delta A &\approx 400 \\
 Z / \Delta Z &\approx 200 \\
 \Delta(\beta\gamma) / \beta\gamma &\approx 5 \cdot 10^{-4}
 \end{aligned}$$

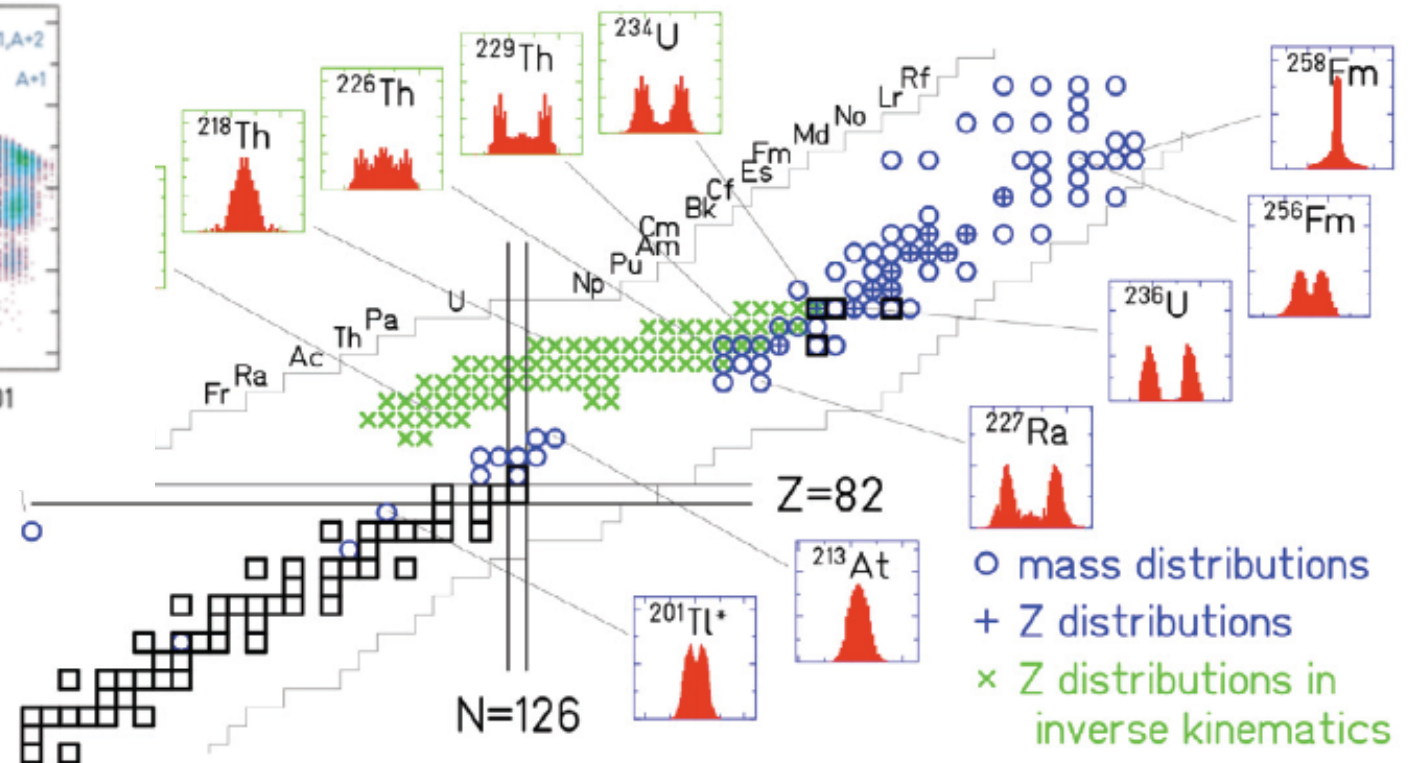
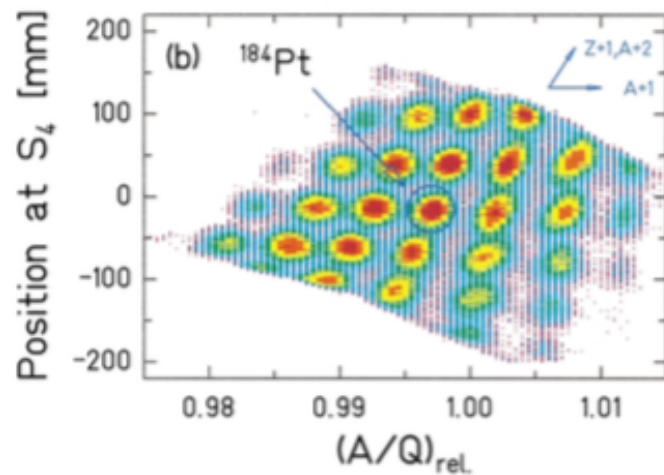
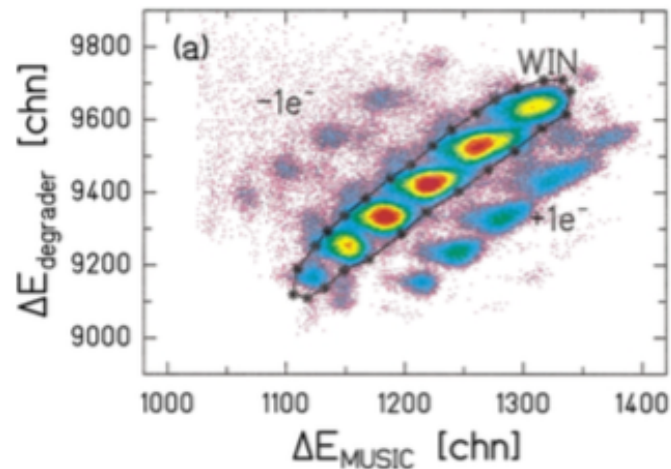


$$\frac{A\beta\gamma}{q} = Br_m$$

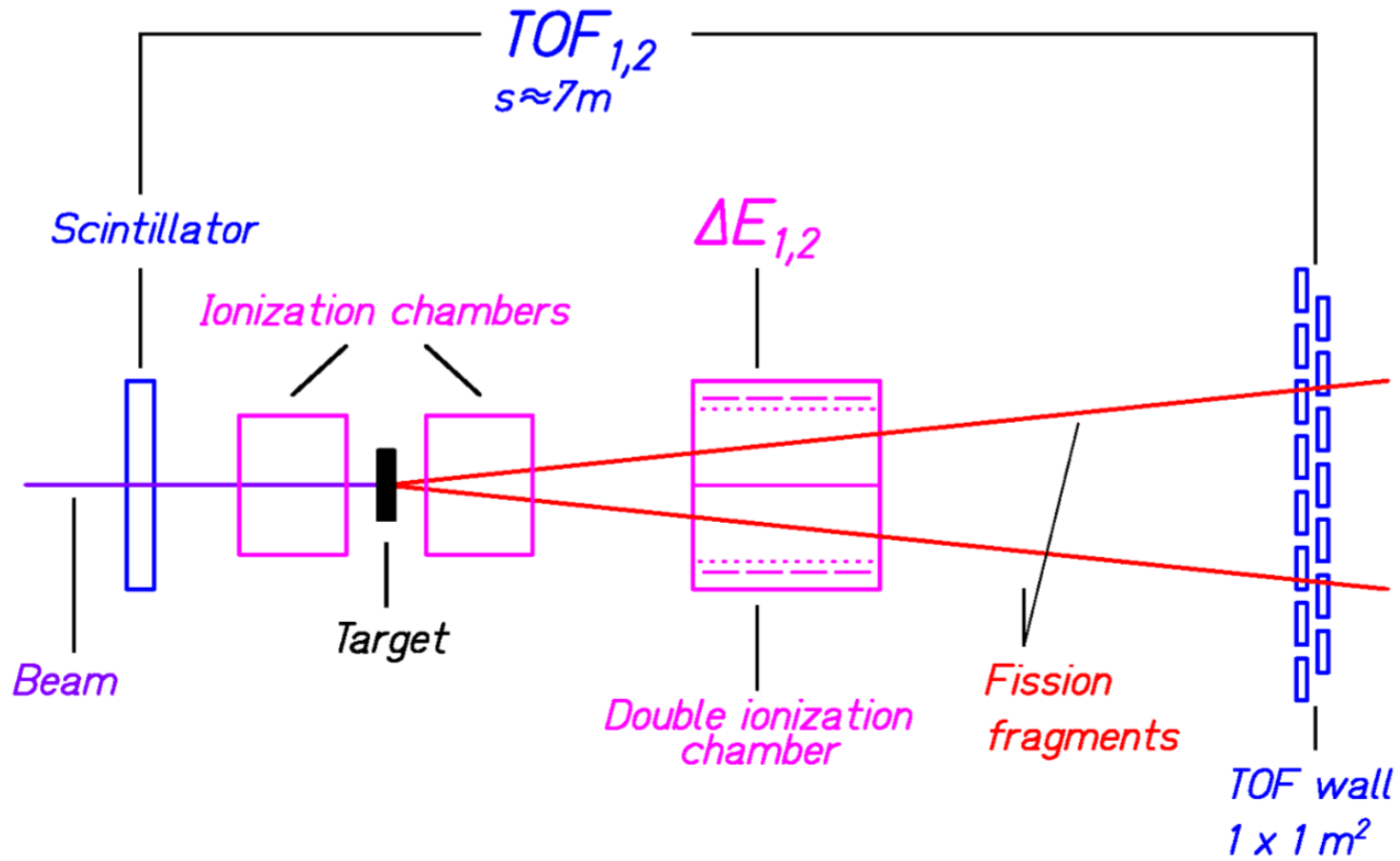
High velocity :  
 $q = Z !!$

# In-flight production of actinides

Large number of fissioning systems !

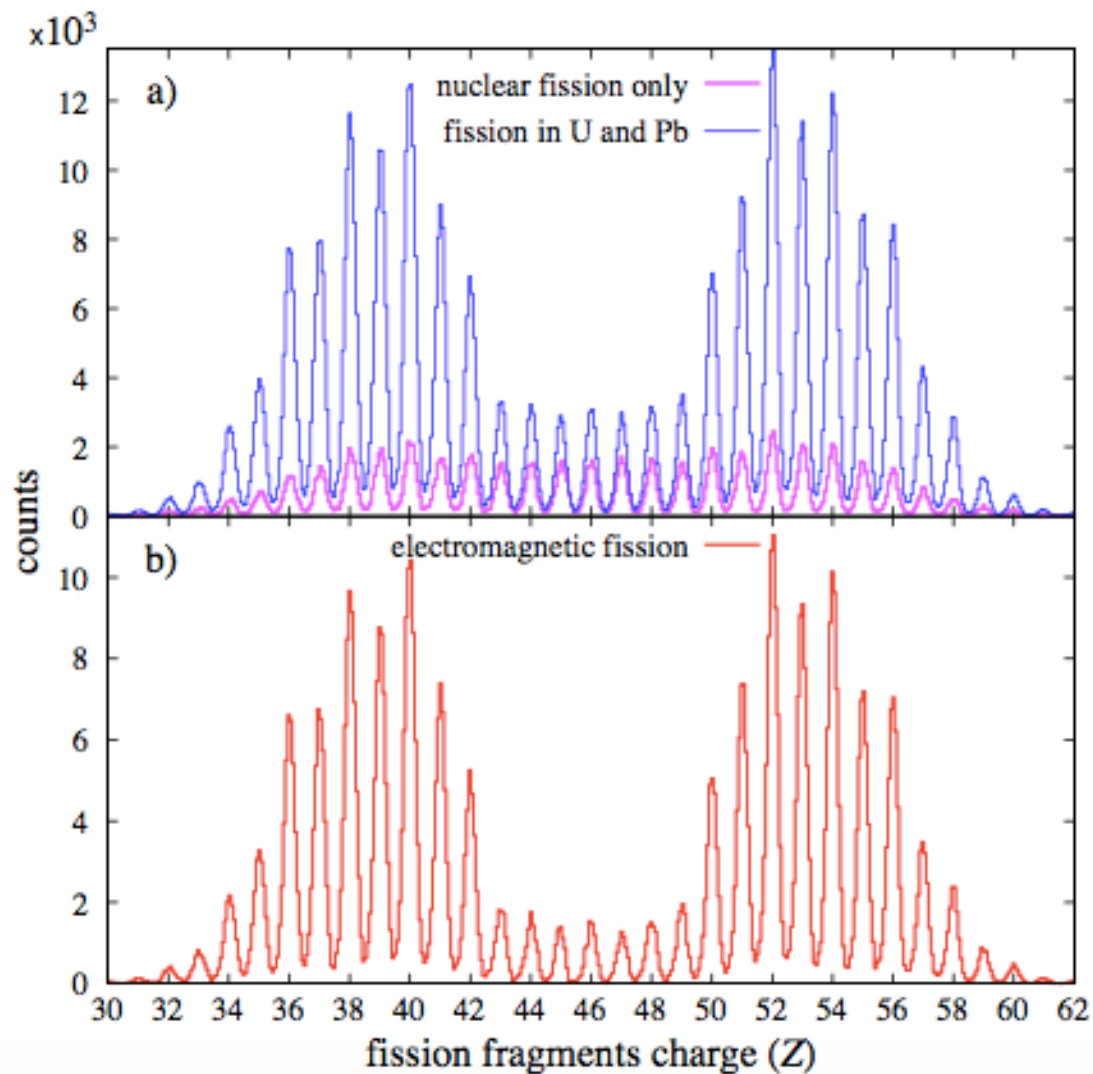


# In-flight fission of actinides



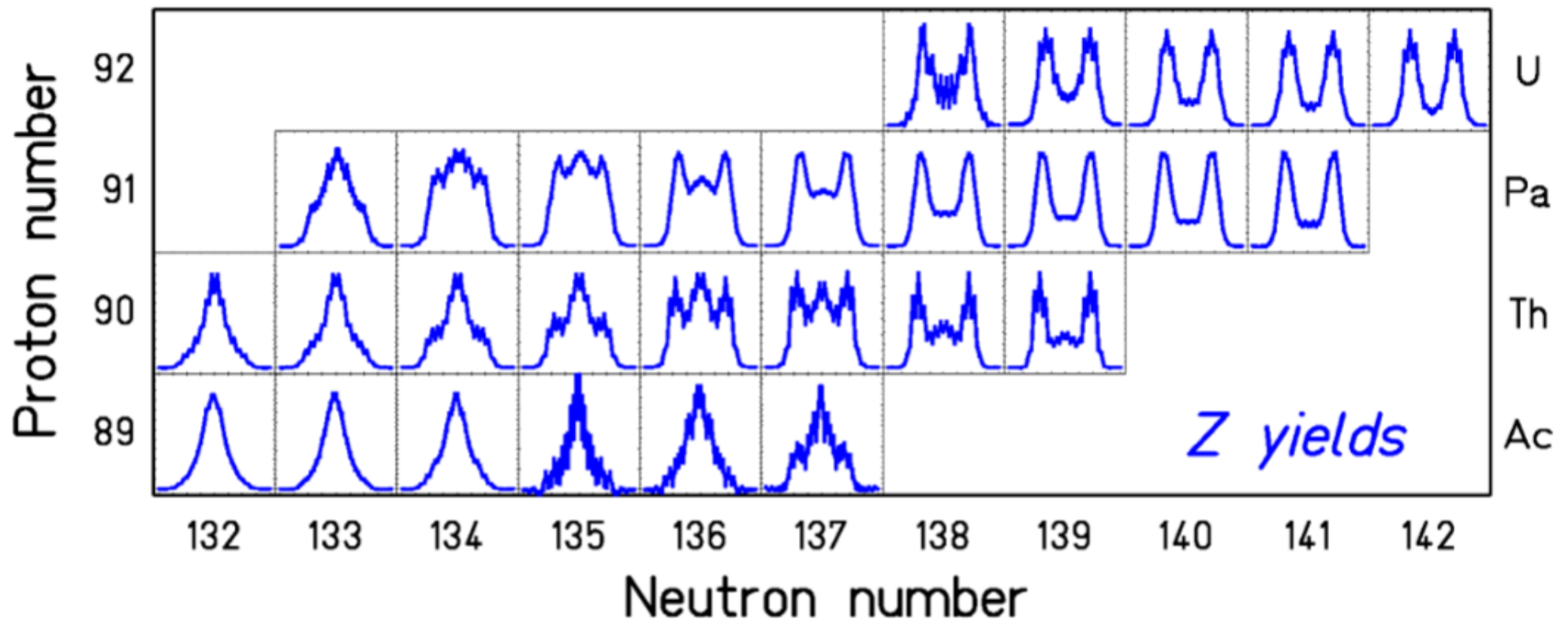


# Improved Z resolution at high kinetic energy



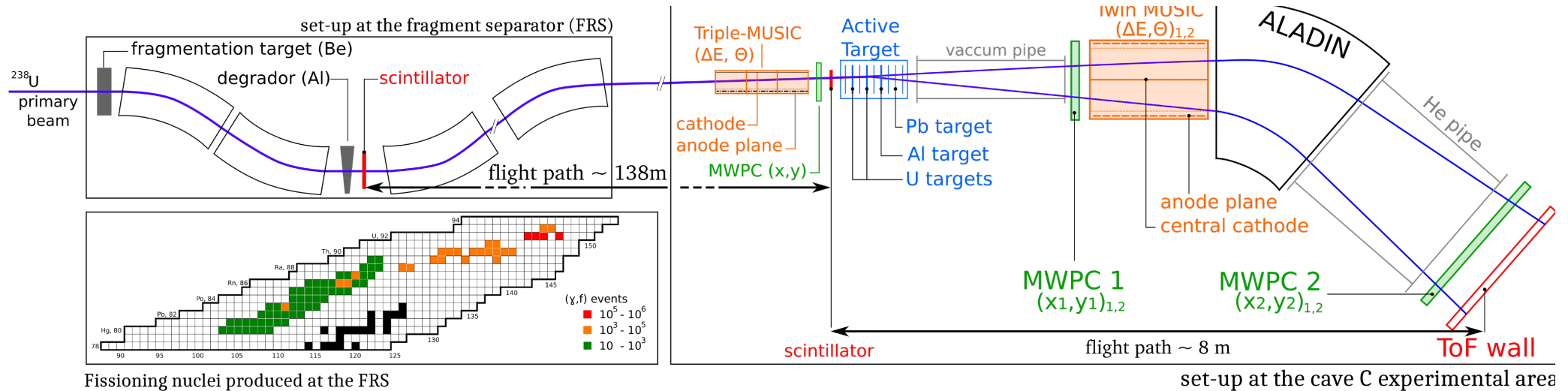
$$\frac{\partial Z}{Z} \sim 0.2\%$$

# Complete distribution of Z yields in many systems



K.-H. Schmidt et al., Nucl. Phys. A 665(2000)

# Isotopic identification at relativistic energy



Fragment separator to produce and select actinides

ALADIN magnet to identify both fission fragments

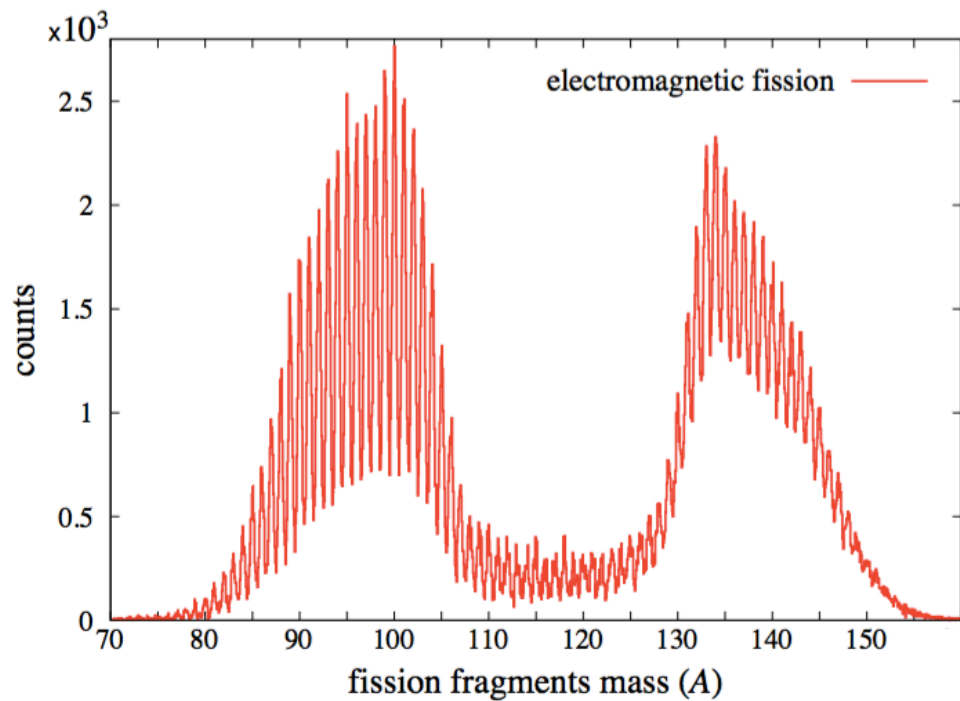
Measure of  $B\rho$ ,  $\Delta E$ , ToF

$$\frac{\partial A}{A} \sim 0.5\%$$

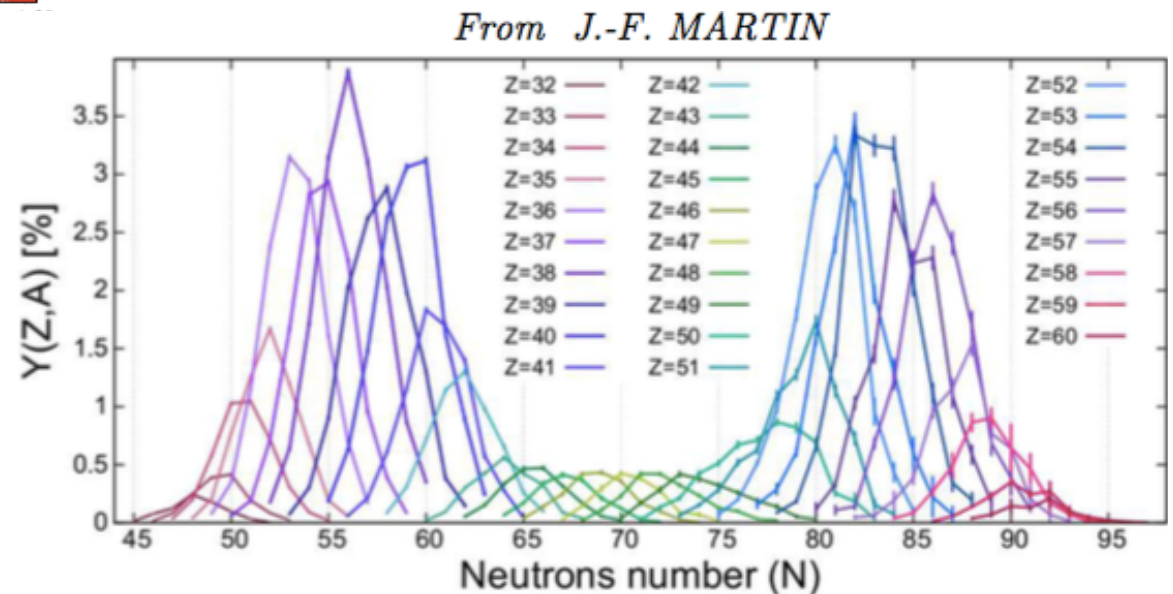
8 m ToF : Absolute challenge to get ToF resolution  
Scintillators with 50ps FWHM resolution

A. Ebran, et al. Nucl. Instr. Meth. A 728 (2013)

# Isotopic identification at relativistic energy



Impressive set of data !!  
Complete isotopic yields  
Very constraining to modelisation !



# Secondary beams

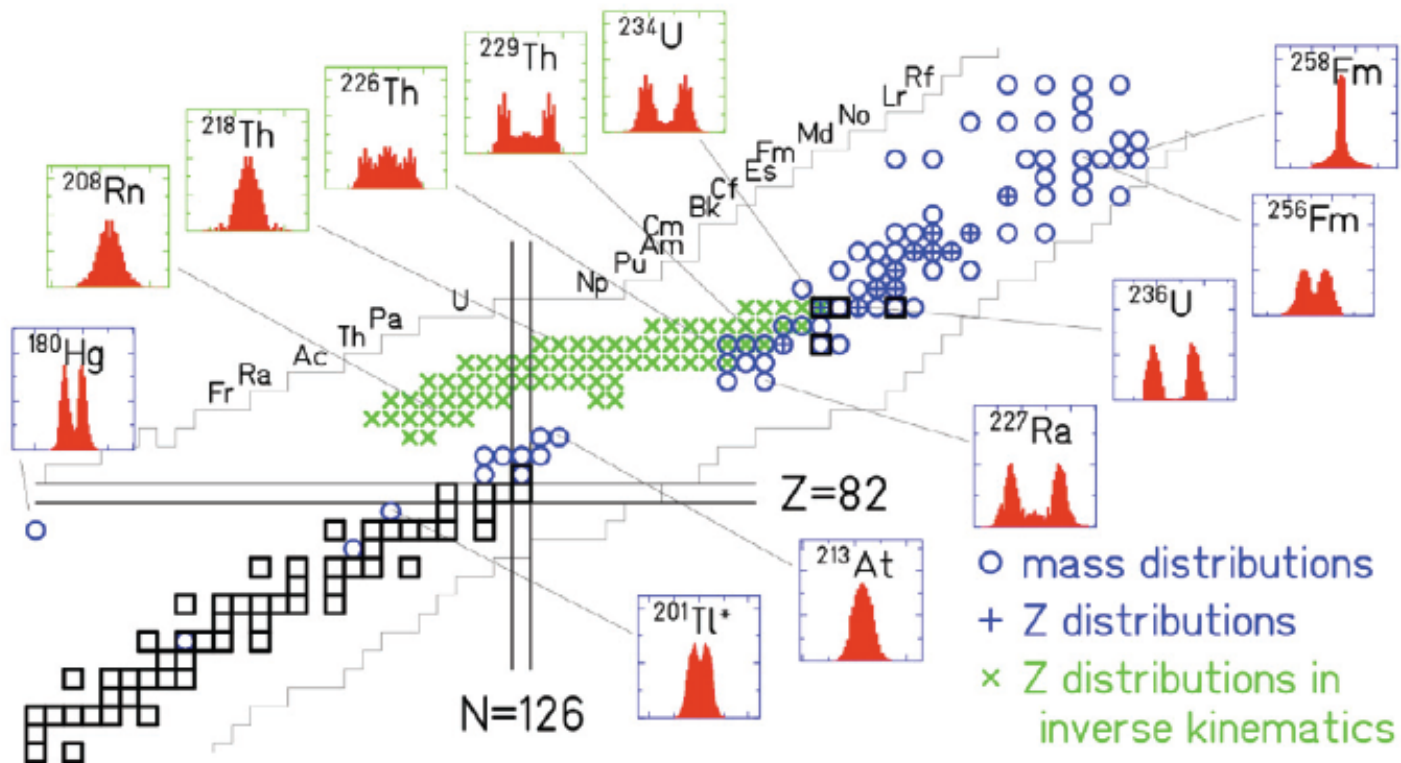
Fission in inverse kinematics

high resolution in  $Z, A$

Complete identification

Both fission-fragments  $\rightarrow$  total neutron multiplicity !

Large number of fissioning systems but light systems



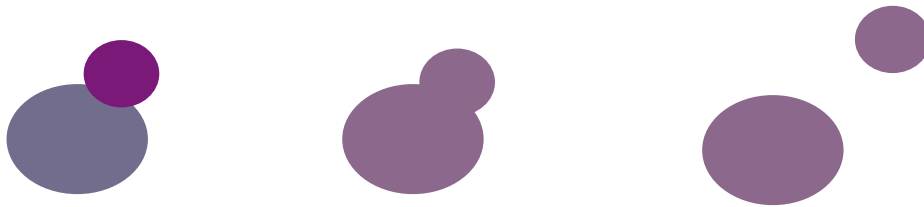
# Multi-nucleon transfer induced fission

Coulomb barrier energy

Very asymmetric reaction :

Nucleon exchange from light to heavy nucleus

Direct reaction

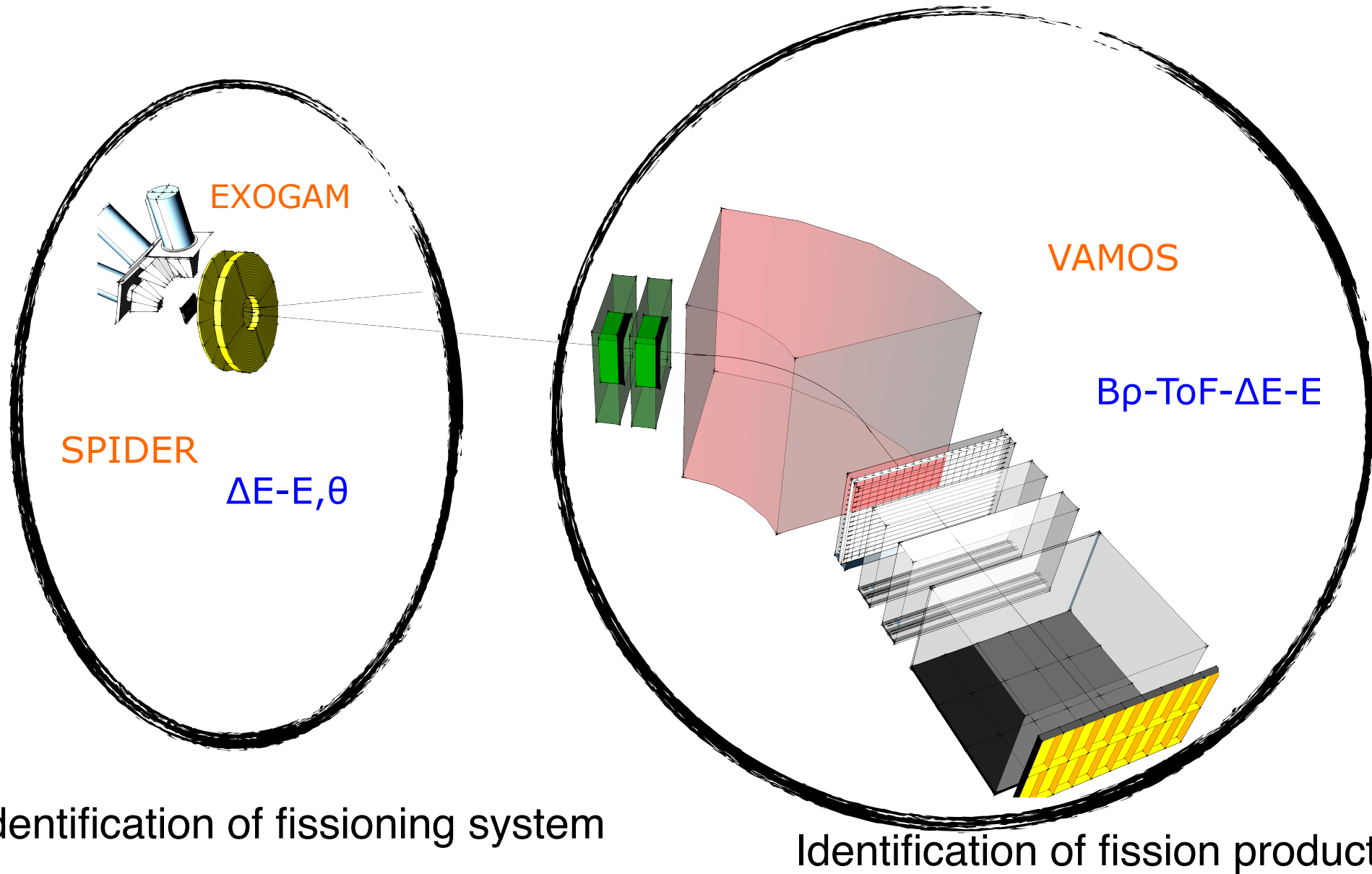


$^{238}\text{U} + ^{12}\text{C}$  :

Production of almost 10  
neutron-rich actinides

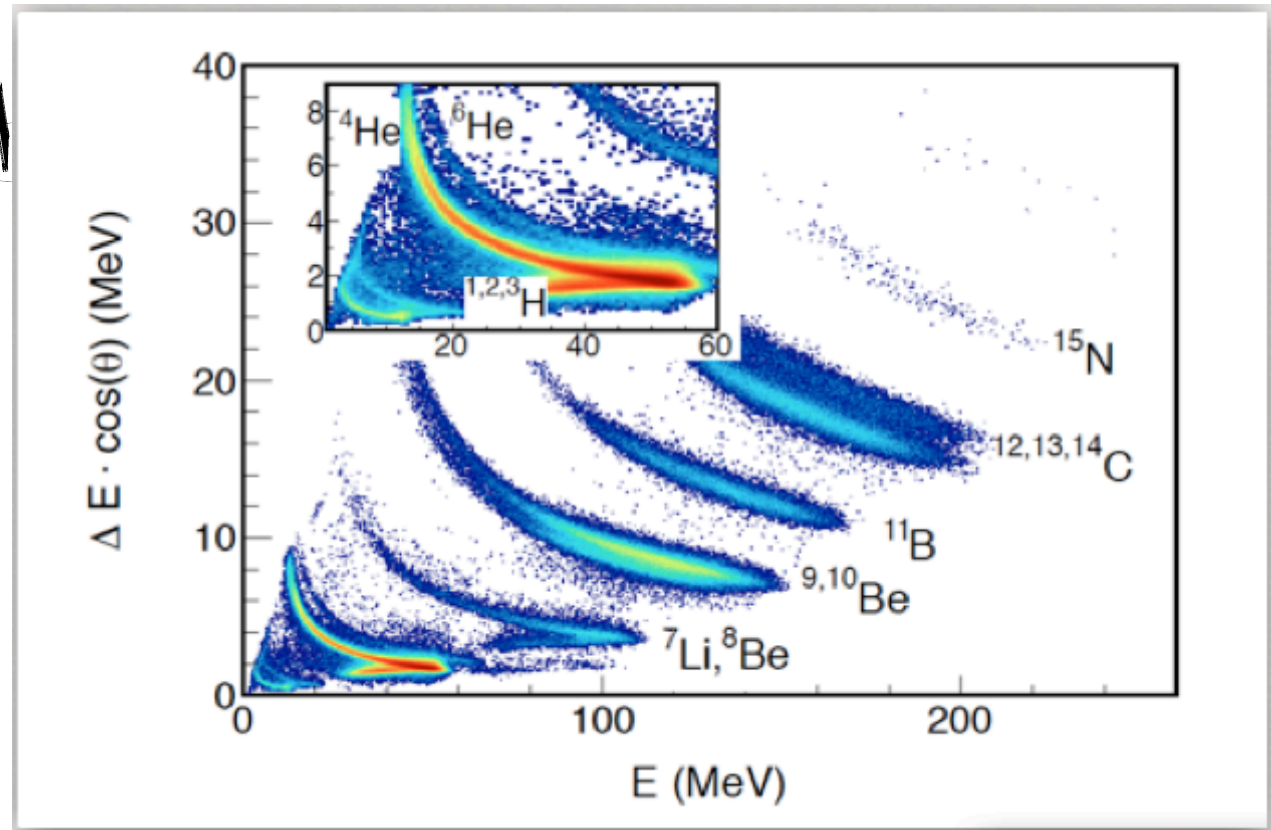
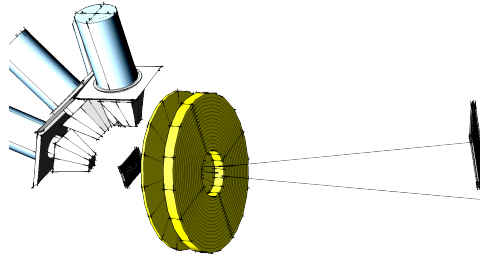
<sup>242</sup> Cf	<sup>243</sup> Cf	<sup>244</sup> Cf	<sup>245</sup> Cf	<sup>246</sup> Cf	<sup>247</sup> Cf	<sup>248</sup> Cf	<sup>249</sup> Cf	<sup>250</sup> Cf	<sup>251</sup> Cf	<sup>252</sup> Cf
<sup>241</sup> Bk	<sup>242</sup> Bk	<sup>243</sup> Bk	<sup>244</sup> Bk	<sup>245</sup> Bk	<sup>246</sup> Bk	<sup>247</sup> Bk	<sup>248</sup> Bk	<sup>249</sup> Bk	<sup>250</sup> Bk	<sup>251</sup> Bk
<sup>240</sup> Cm	<sup>241</sup> Cm	<sup>242</sup> Cm	<sup>243</sup> Cm	<sup>244</sup> Cm	<sup>245</sup> Cm	<sup>246</sup> Cm	<sup>247</sup> Cm	<sup>248</sup> Cm	<sup>249</sup> Cm	<sup>250</sup> Cm
<sup>239</sup> Am	<sup>240</sup> Am	<sup>241</sup> Am	<sup>242</sup> Am	<sup>243</sup> Am	<sup>244</sup> Am	<sup>245</sup> Am	<sup>246</sup> Am	<sup>247</sup> Am	<sup>248</sup> Am	<sup>249</sup> Am
<sup>238</sup> Pu	<sup>239</sup> Pu	<sup>240</sup> Pu	<sup>241</sup> Pu	<sup>242</sup> Pu	<sup>243</sup> Pu	<sup>244</sup> Pu	<sup>245</sup> Pu	<sup>246</sup> Pu	<sup>247</sup> Pu	
<sup>237</sup> Np	<sup>238</sup> Np	<sup>239</sup> Np	<sup>240</sup> Np	<sup>241</sup> Np	<sup>242</sup> Np	<sup>243</sup> Np	<sup>244</sup> Np			
<sup>236</sup> U	<sup>237</sup> U	<sup>238</sup> U	<sup>239</sup> U	<sup>240</sup> U	<sup>241</sup> U	<sup>242</sup> U				

# Multi-nucleon transfer induced fission





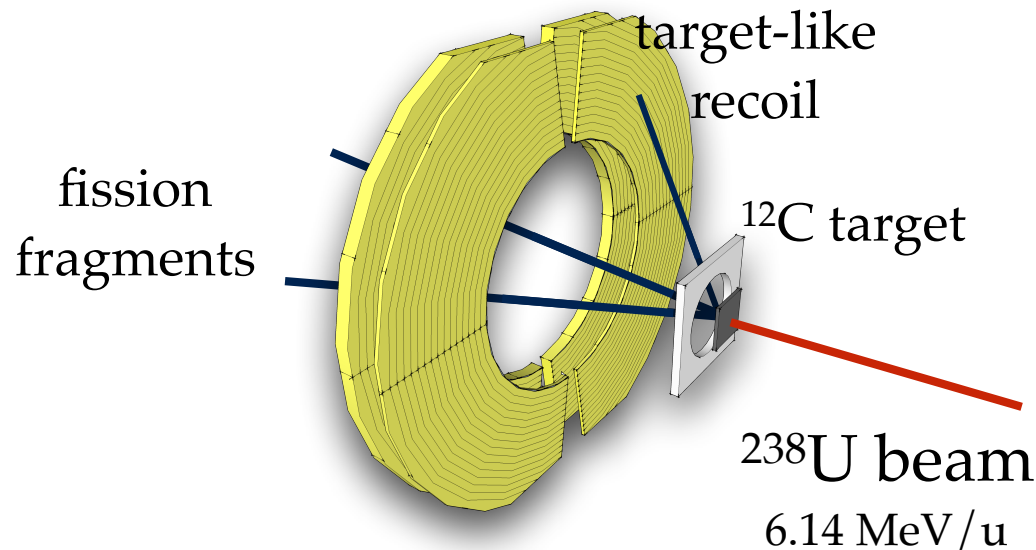
# Identification of the transfer channels



C. Rodriguez-Tajes et al.,  
 PRC89 (2014) 024614

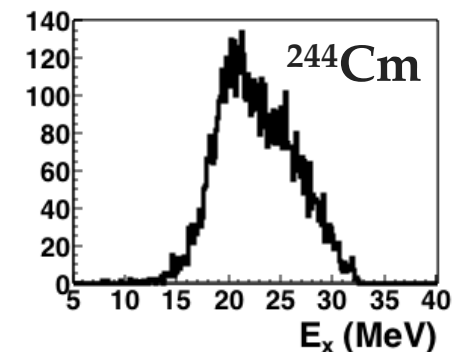
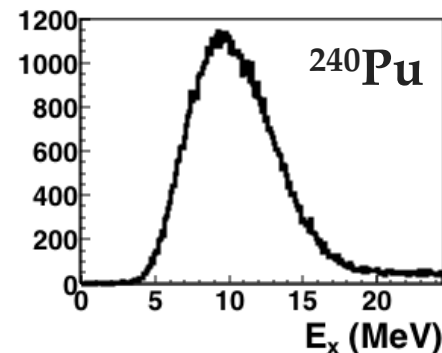
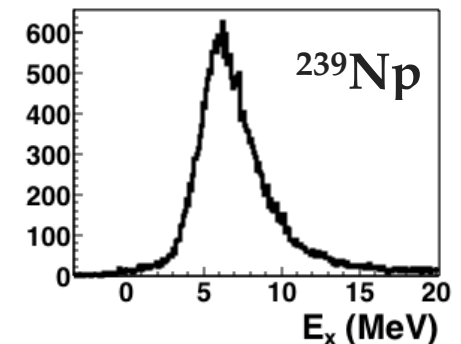
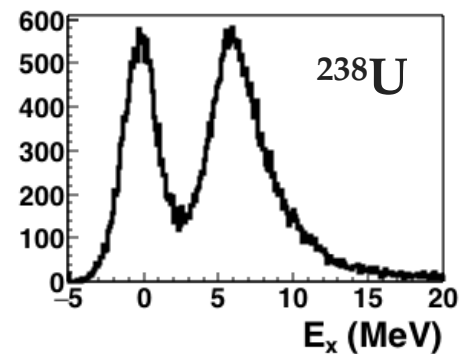


# Identification of the transfer channels

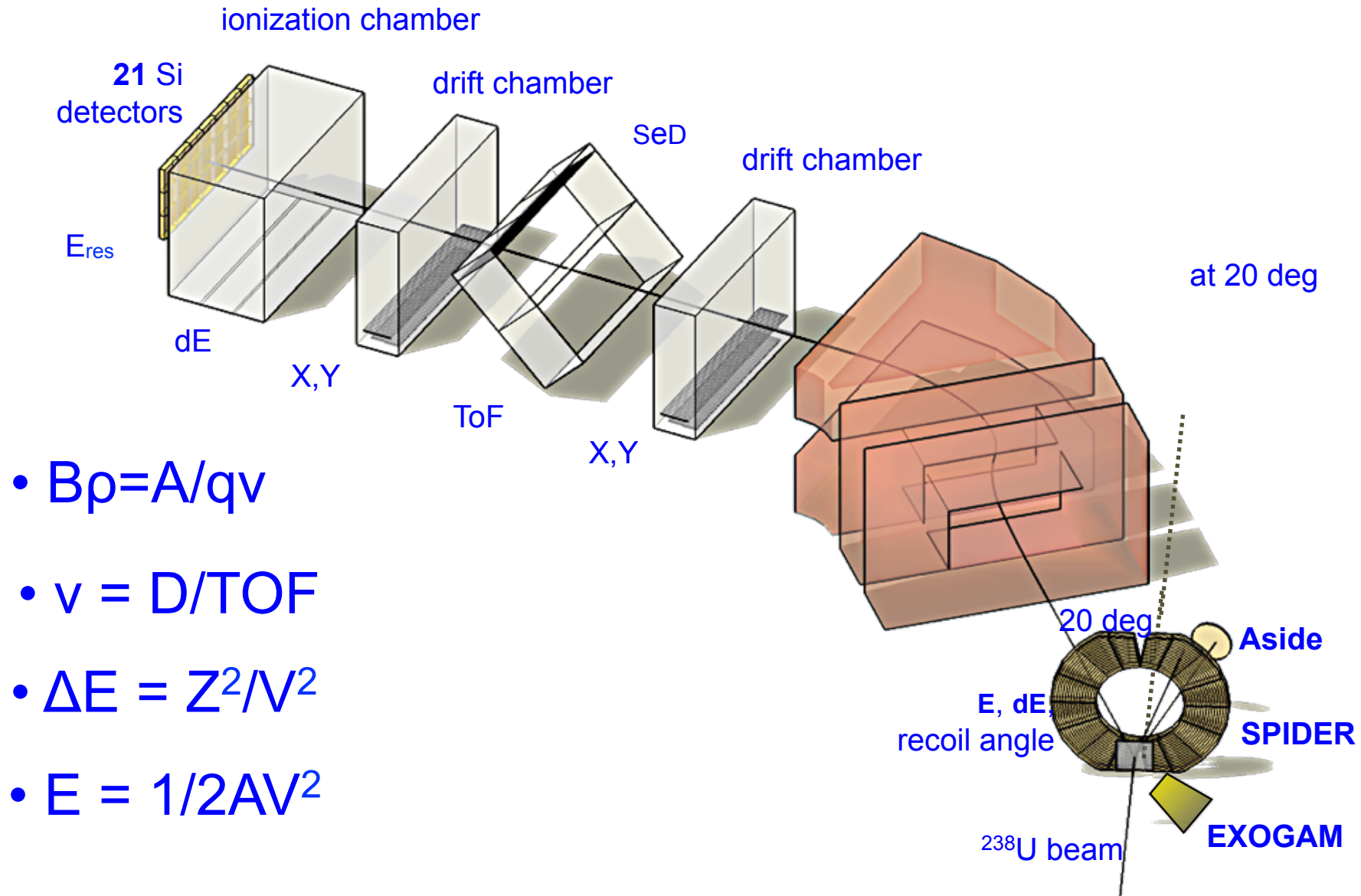


$^{238}\text{U} (^{12}\text{C}, ^{12}\text{C}) ^{238}\text{U}$   
 $^{238}\text{U} (^{12}\text{C}, ^{11}\text{B}) ^{239}\text{Np}$   
 $^{238}\text{U} (^{12}\text{C}, ^{10}\text{Be}) ^{240}\text{Pu}$   
 $^{238}\text{U} (^{12}\text{C}, ^6\text{He}) ^{244}\text{Cm}$

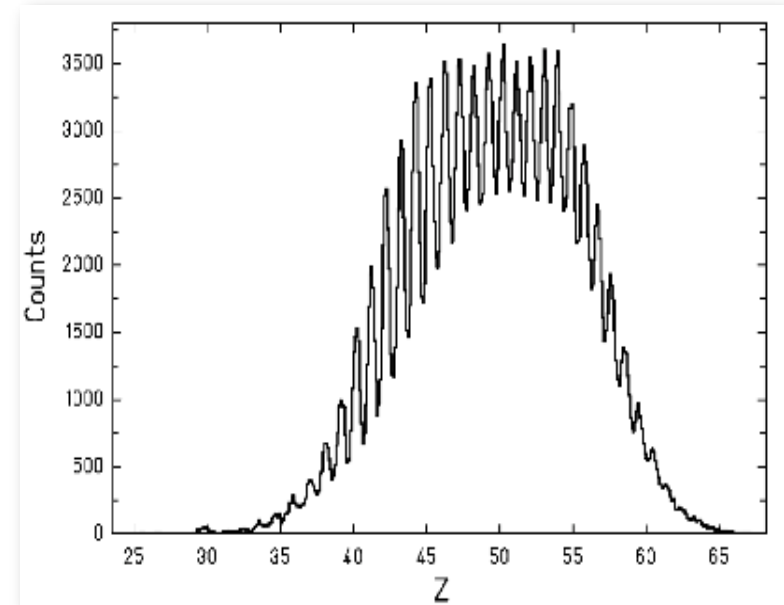
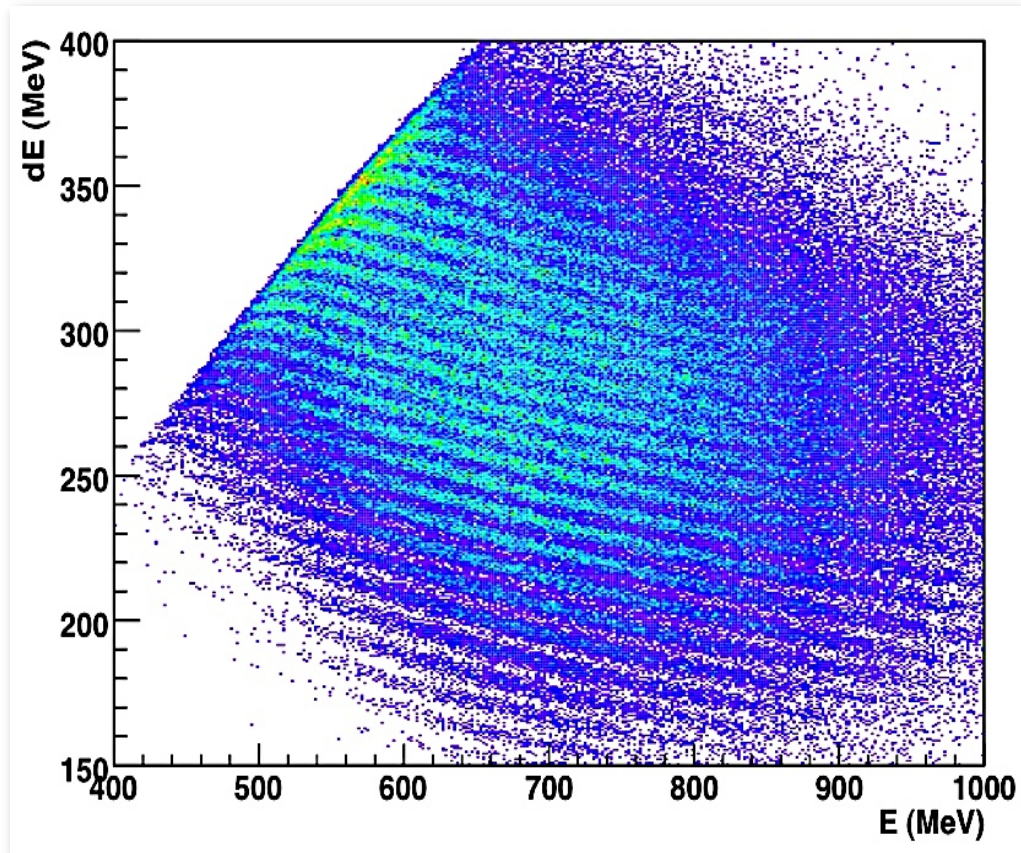
- Isotopic identification
- Reconstruction of binary reaction
- $E_x$  reconstruction



# Identification of fission products



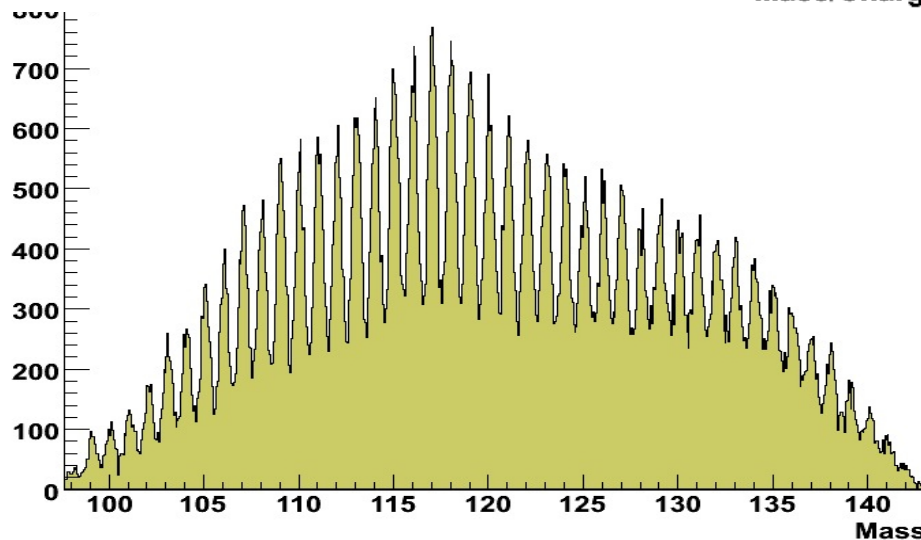
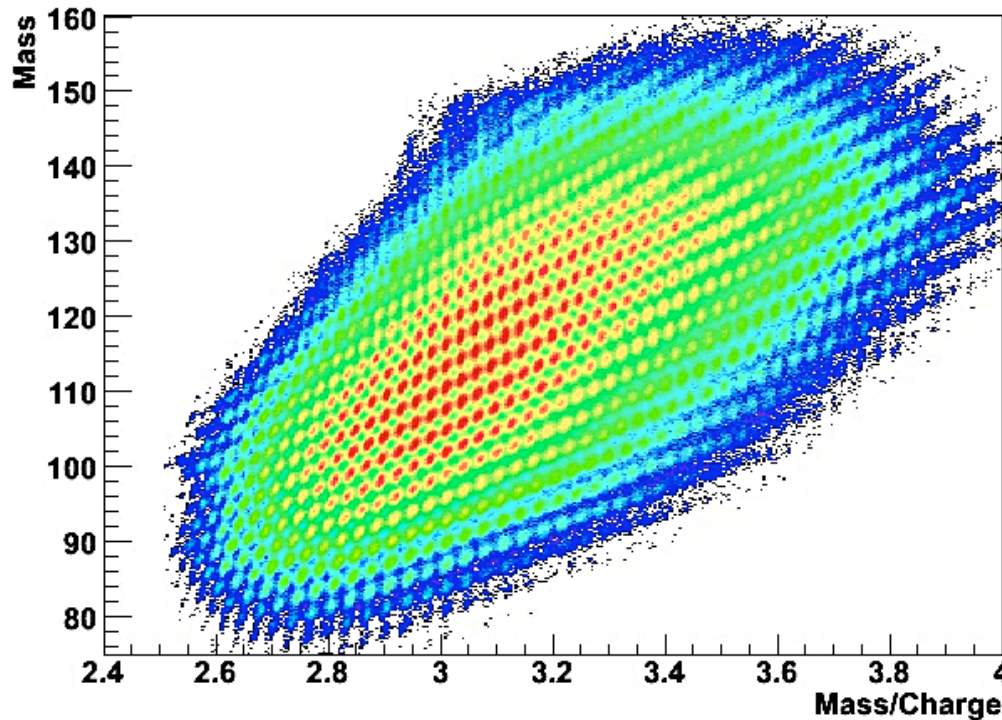
# Improved atomic number resolution



$$\Delta Z/Z \approx 1.5 \cdot 10^{-2}$$



# Mass resolution through magnetic rigidity resolution

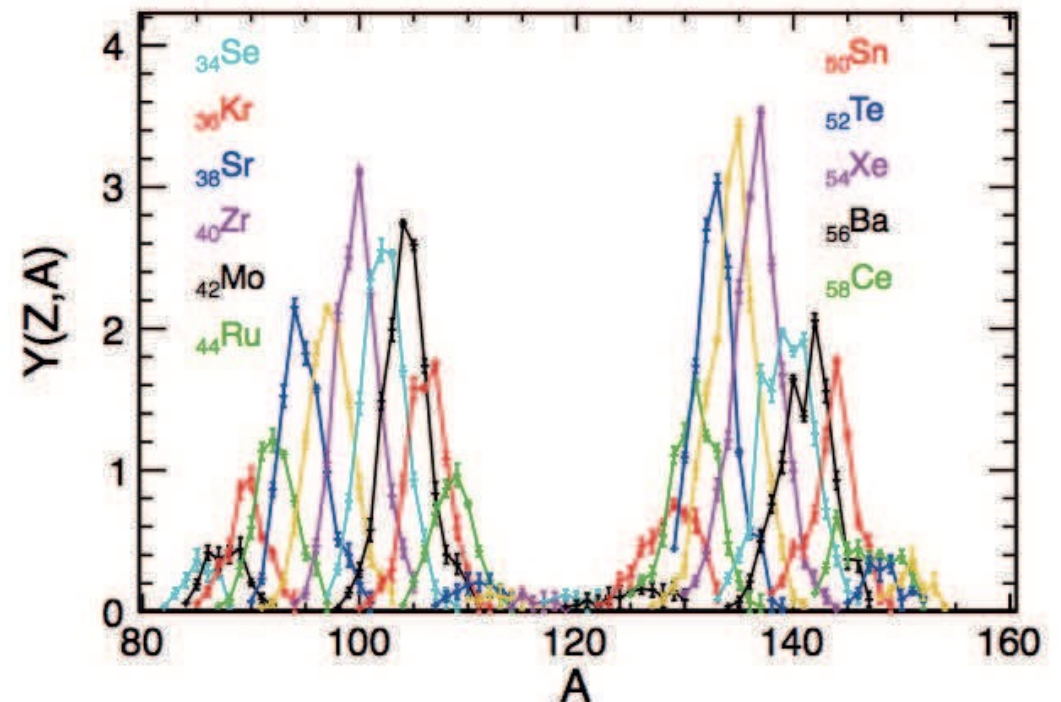


$$A = E/(\gamma - 1)$$

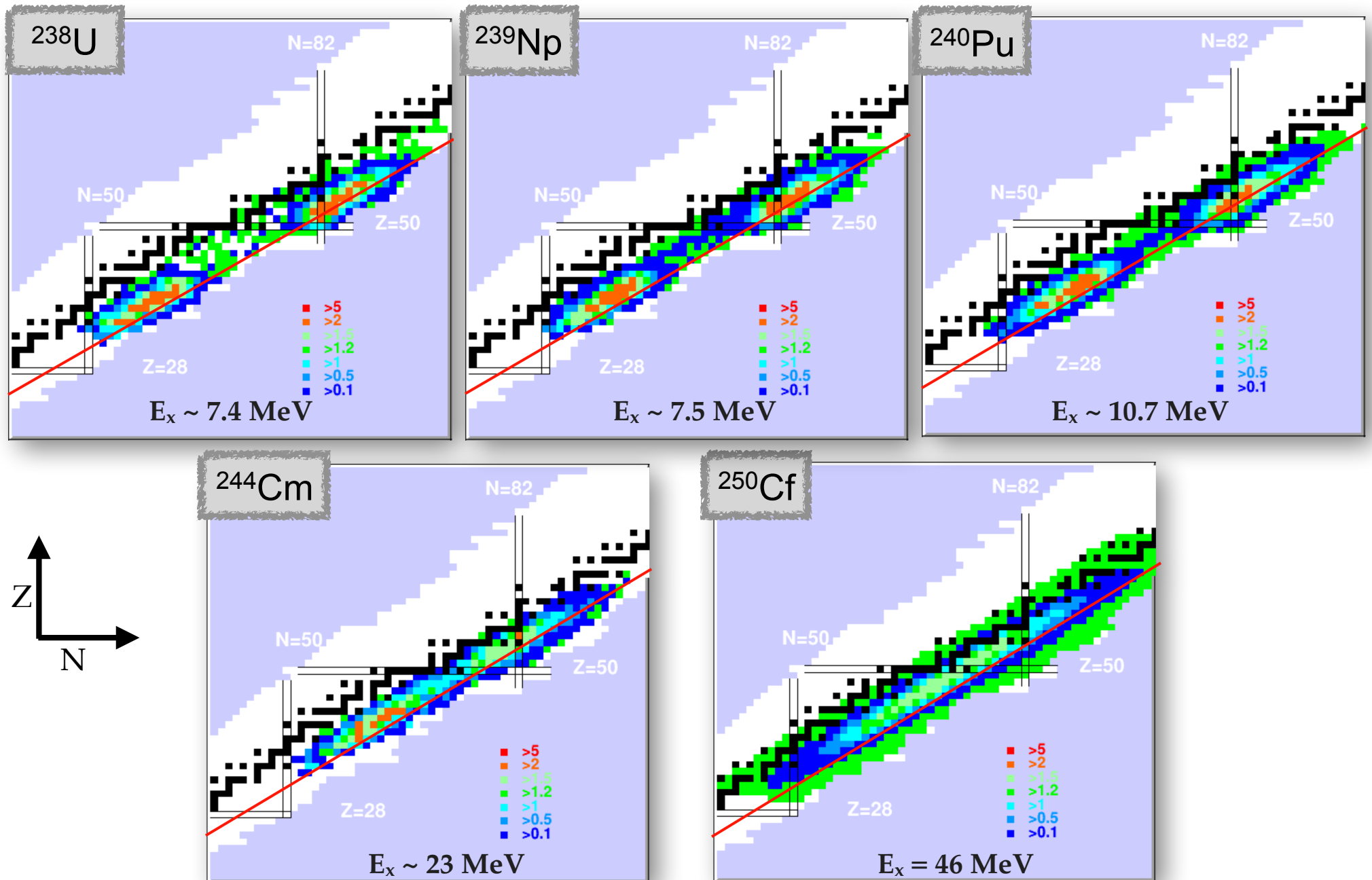
$$A/q = B\rho/(\beta\gamma)$$

$$dq/q \sim 1.5 \%$$

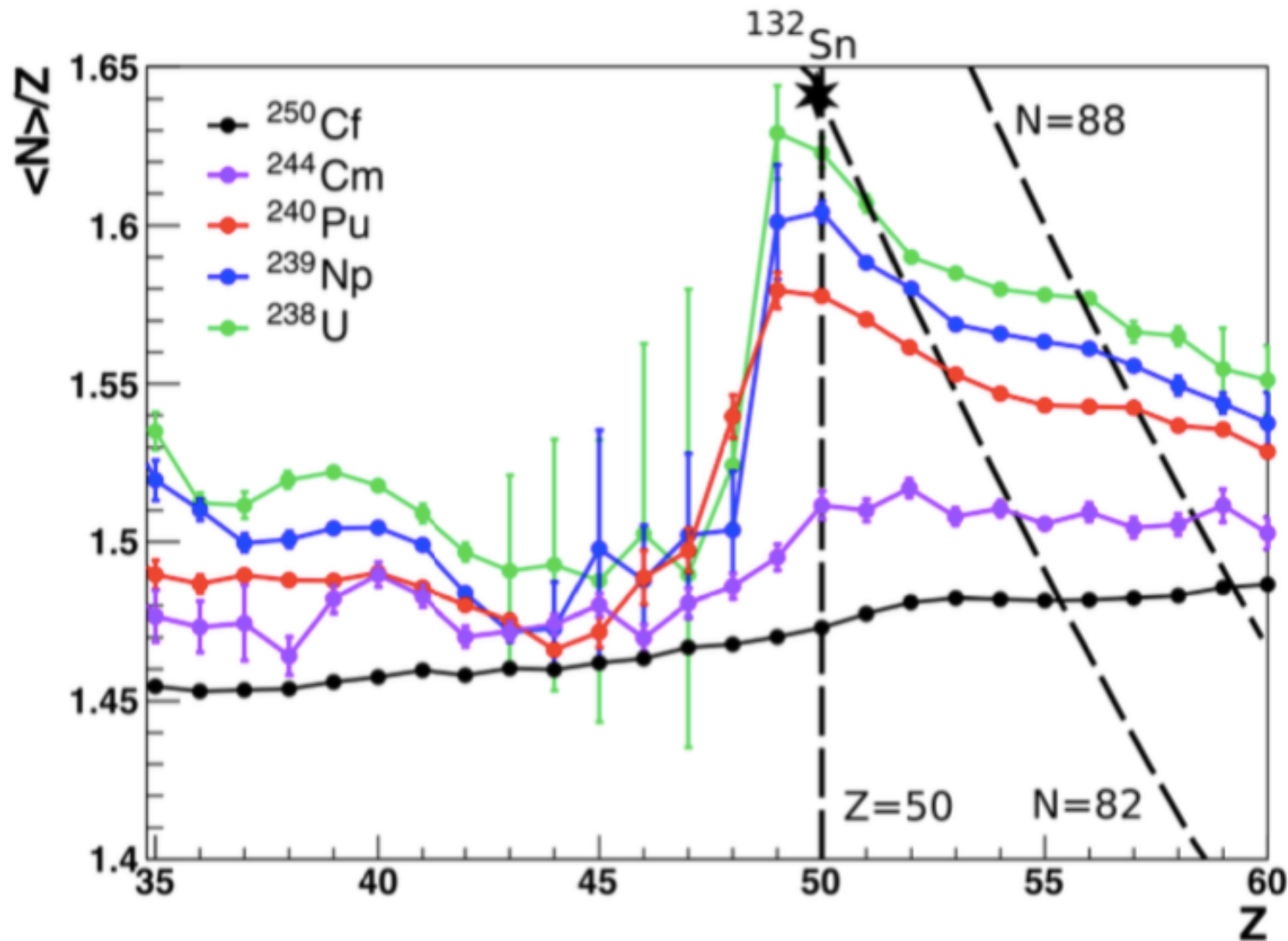
$$dA/A \sim 0.6 \%$$



# Results: Isotopic Fission Yields



New data : new type of observables :  
 Charge polarisation



**New techniques allow for improved resolution for the identification of fission fragments**

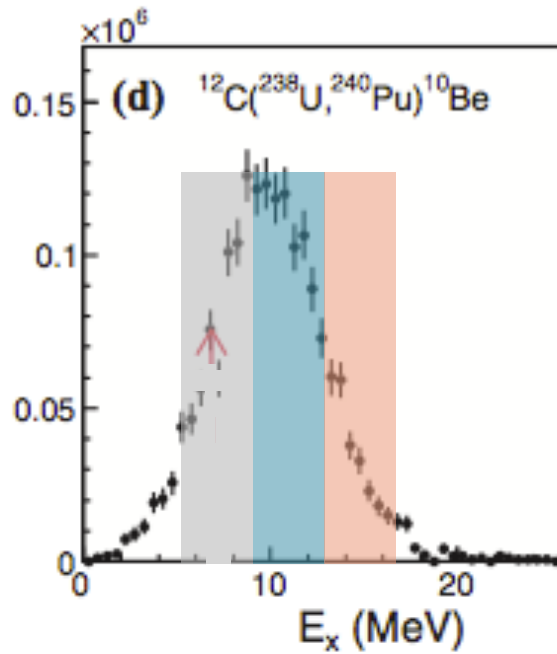
**In addition they allow for :**

**2D-map of the fission fragments**

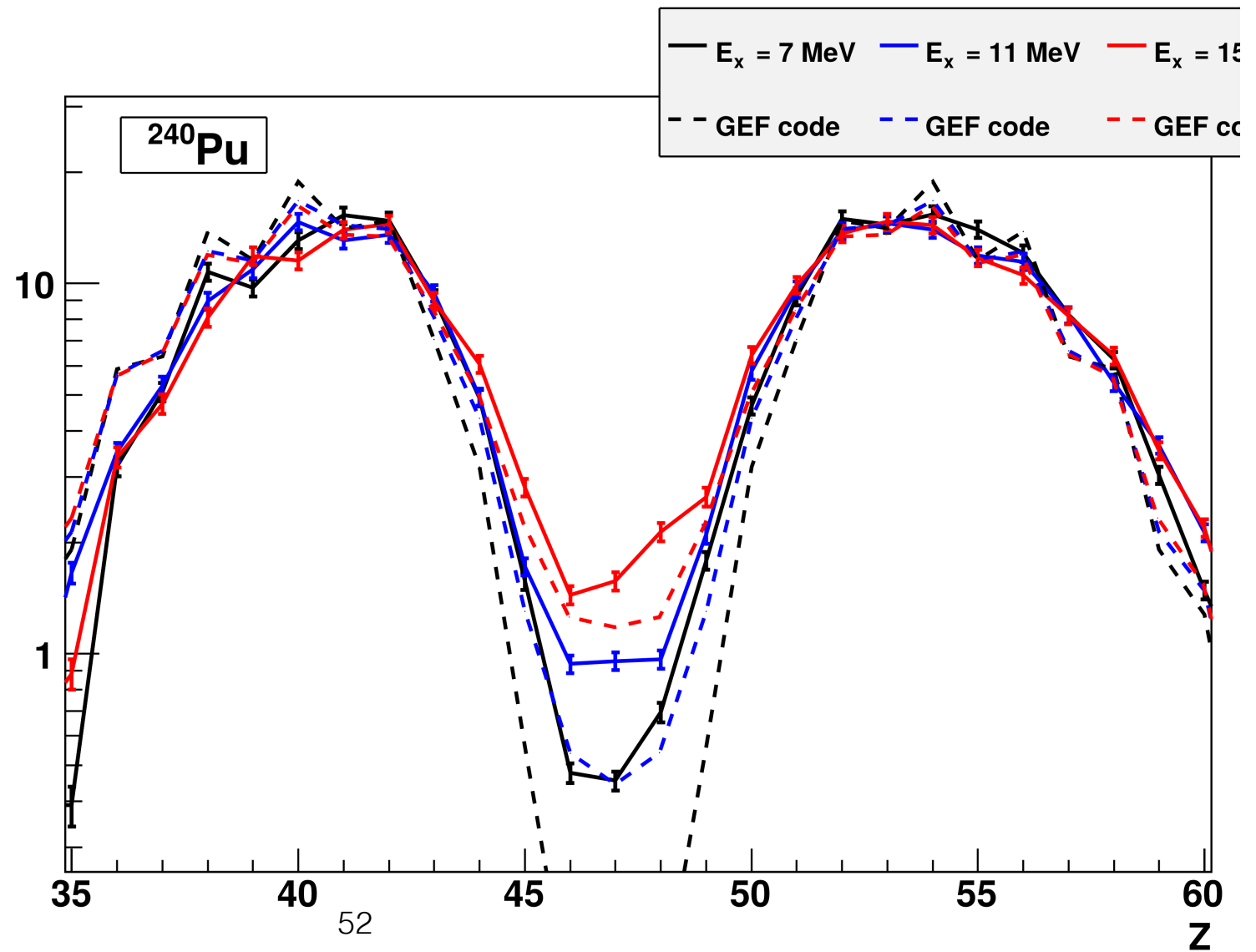
**study the evolution with excitation energy**

**improving the number of fissioning actinides due to nuclear reactions**

# Evolution of mass distribution with Excitation energy

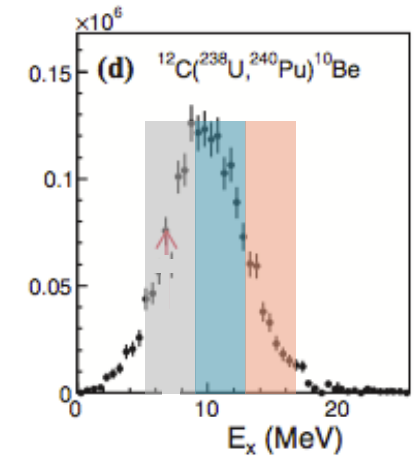
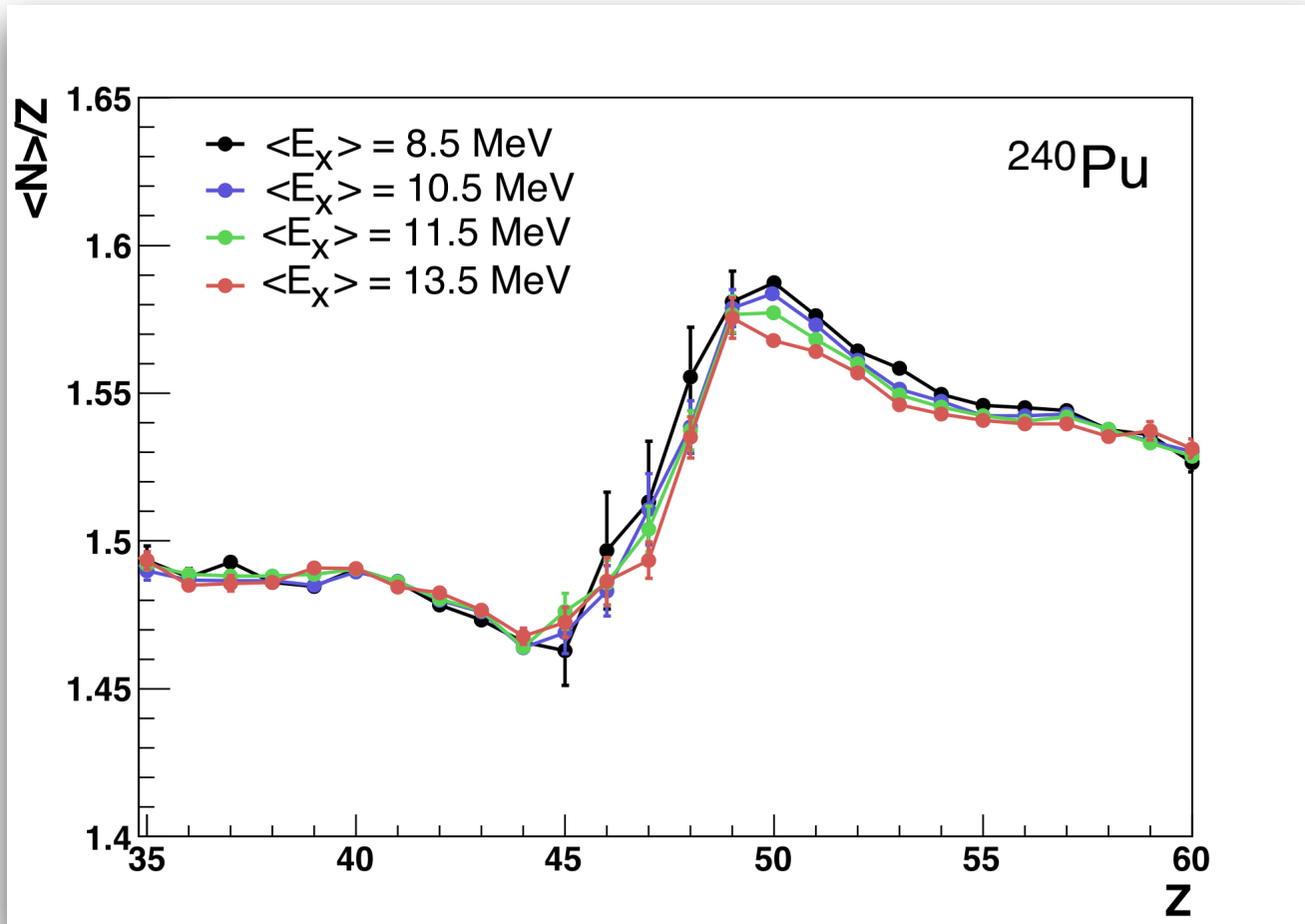


From transfer kinematics



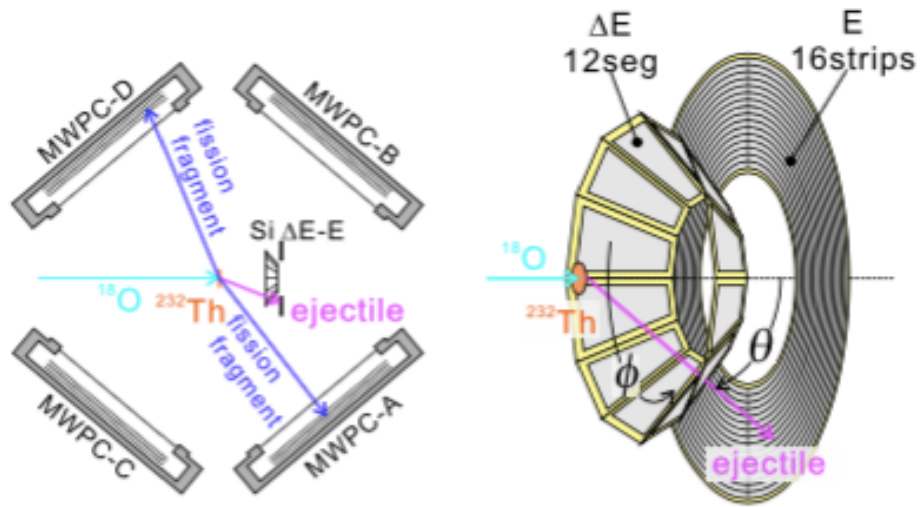


# Evolution of charge polarisation with Excitation energy

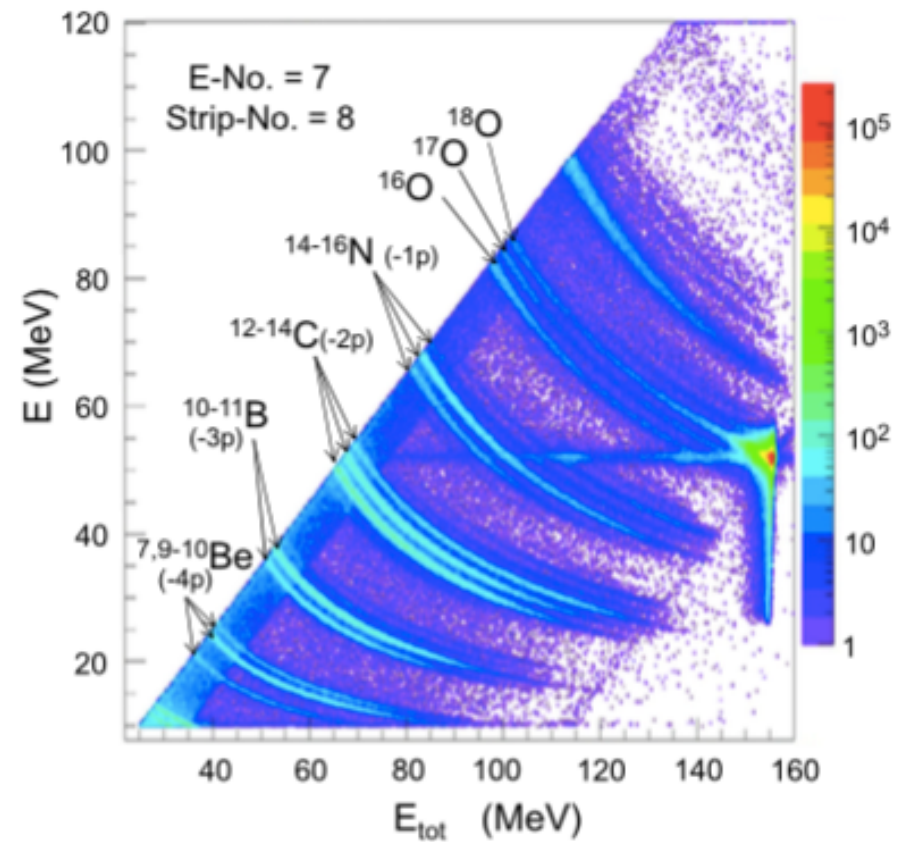


Only heavy fragments evaporate neutrons !!

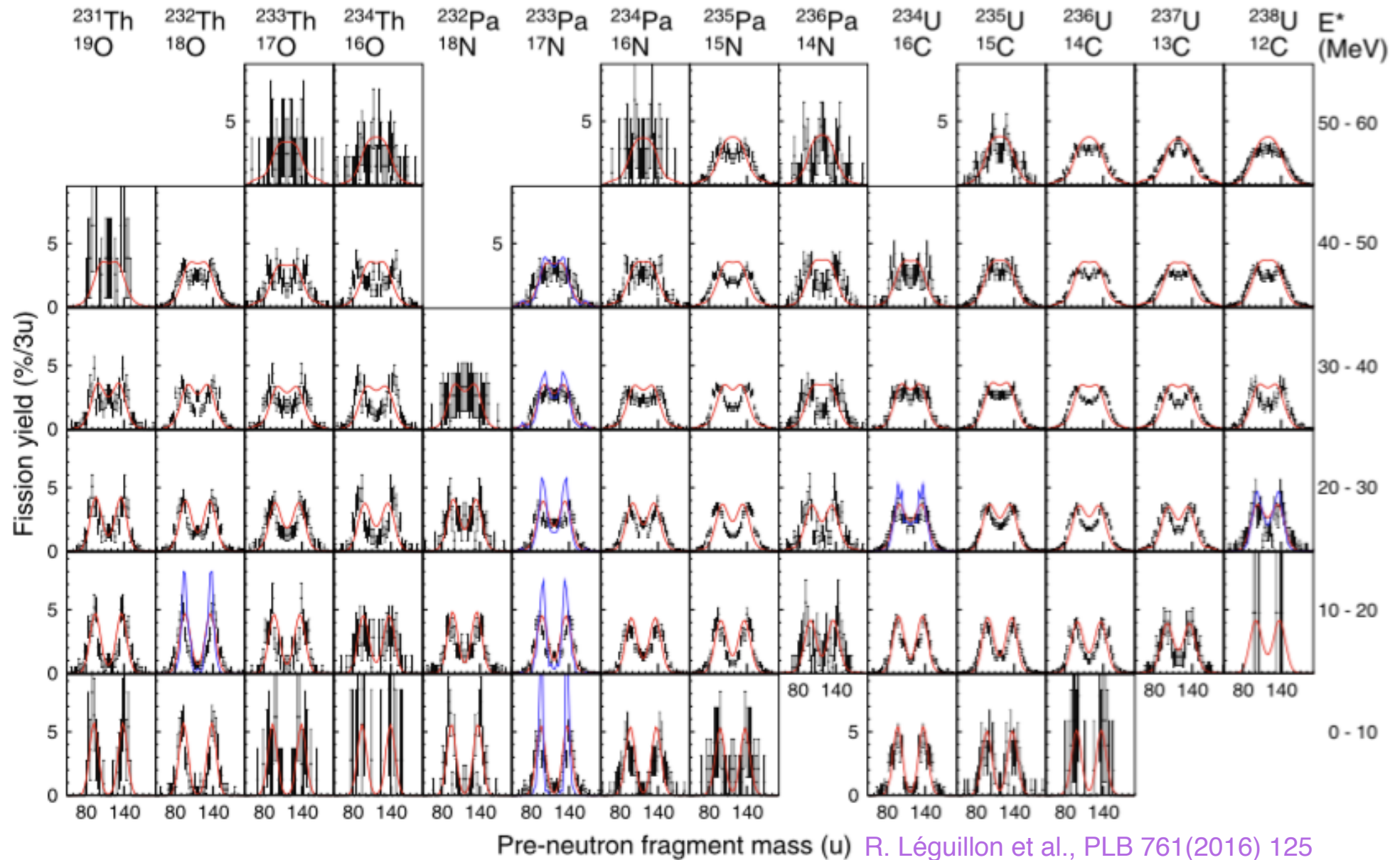
# Multi-nucleon reaction used in direct kinematics



K. Nishio,, JAEA Tandem



# Multi-nucleon reaction used in direct kinematics



## 7. Fission Yields : systematic uncertainties and errors

Fission yields : number of fragment per fission event

$$\sum Y(Z, A) = 200$$

2 fission fragments/ fission

$$Y(Z, A) = \frac{\partial N_f}{\partial Z \partial A}$$

isotopic identification  
 gamma spectroscopy,  
 energy-loss, energy and time-of flight

$$Y(A) = \sum_Z Y(Z, A) = \frac{\partial N_f}{\partial A}$$

mass identification  
 energy, time-of-flight measurements

$$Y(Z) = \sum_A Y(Z, A) = \frac{\partial N_f}{\partial Z}$$

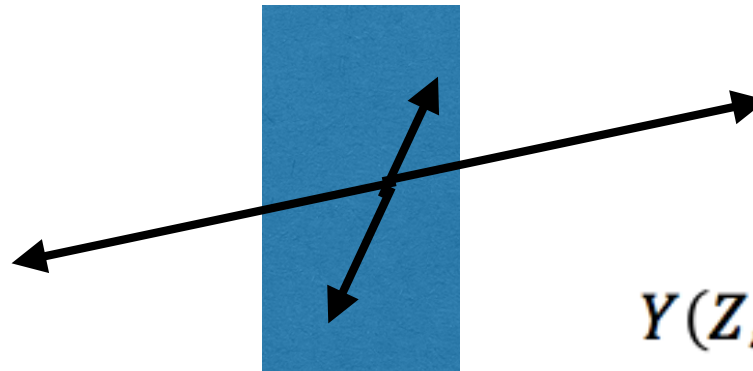
atomic number identification  
 energy-loss, X-ray

## 7. Fission Yields : systematic uncertainties and errors

Relative yields : beam intensity, target thickness, .. are not key parameters

Detection relative efficiency :

- — -from  $Z \sim 30$  to  $Z \sim 60$ , energy-loss varies with a factor 4
- — target release efficiency : depends on  $Z$ , and angle

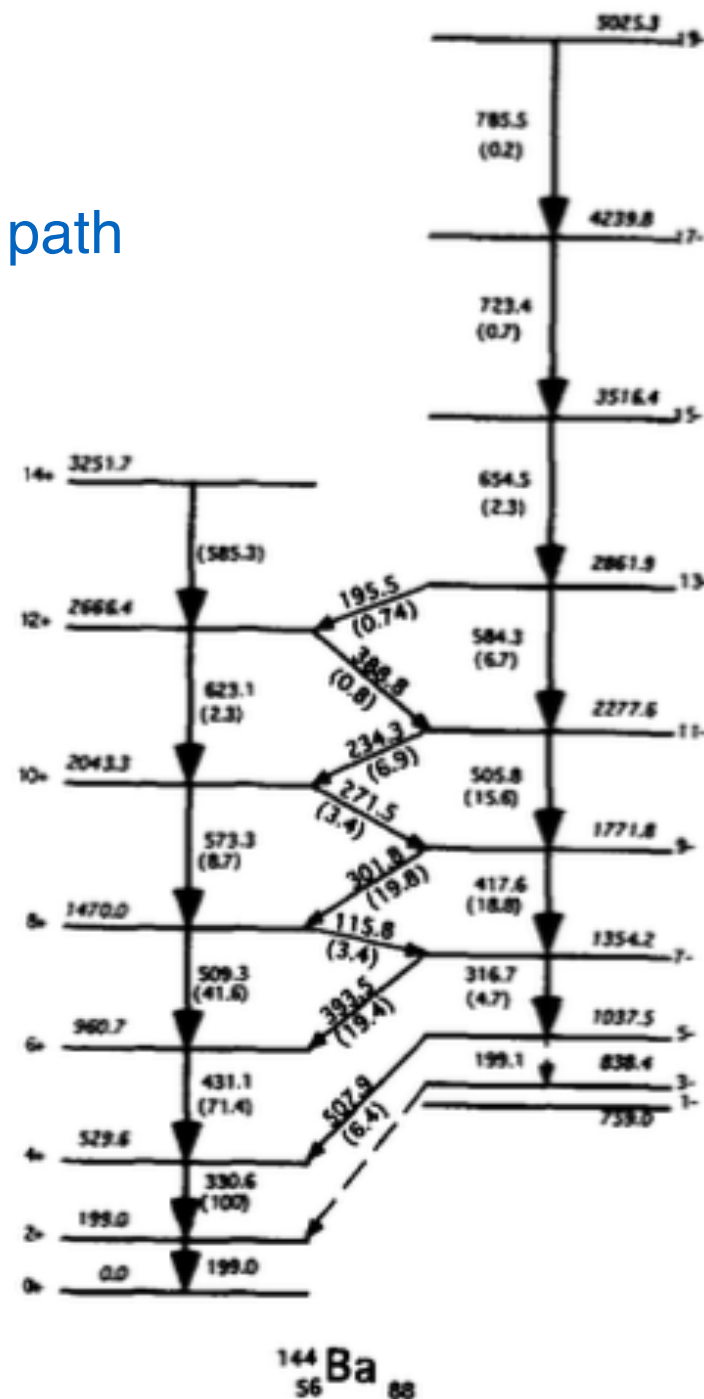
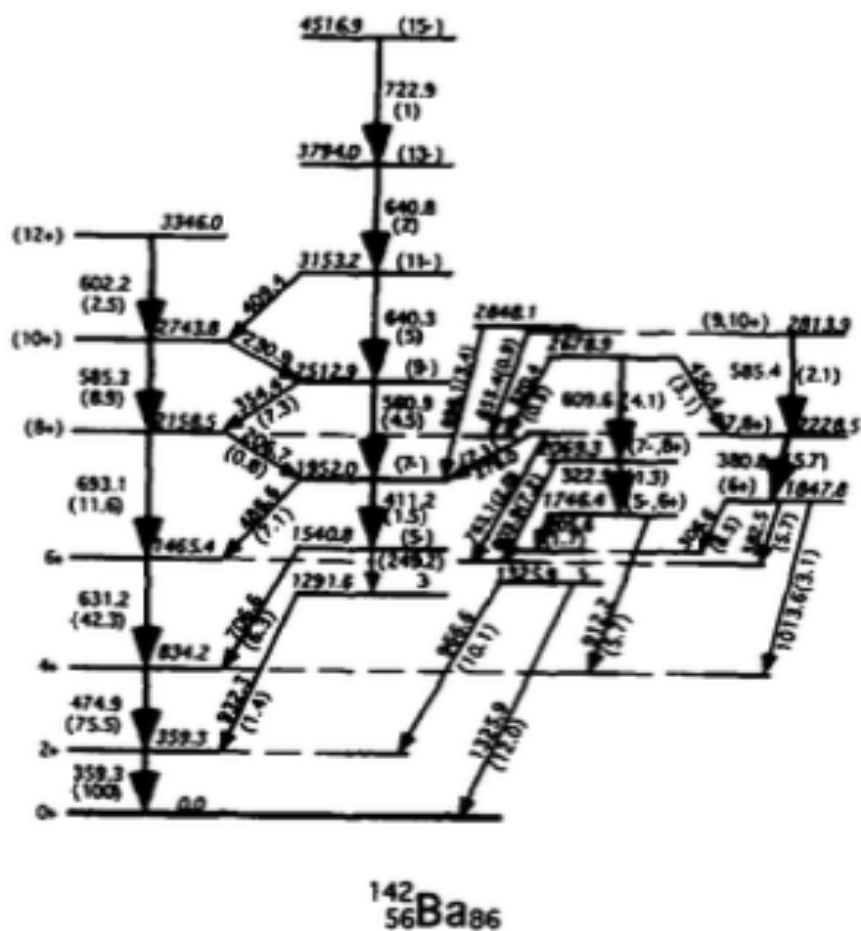


$$Y(Z, A) = \frac{\partial N_f}{\partial Z \partial A} \frac{1}{\xi(Z, A, E)}$$

Need a precise simulation of the detection set-up

## 7. Fission Yields : systematic uncertainties and errors

Uncertainty on level scheme:  
isomeric states ; fragmentation of the decay path

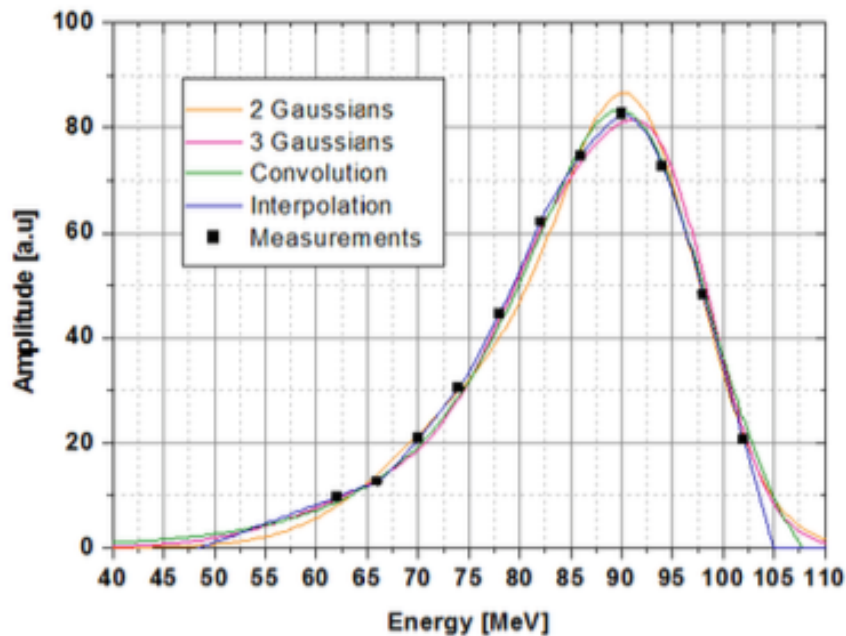
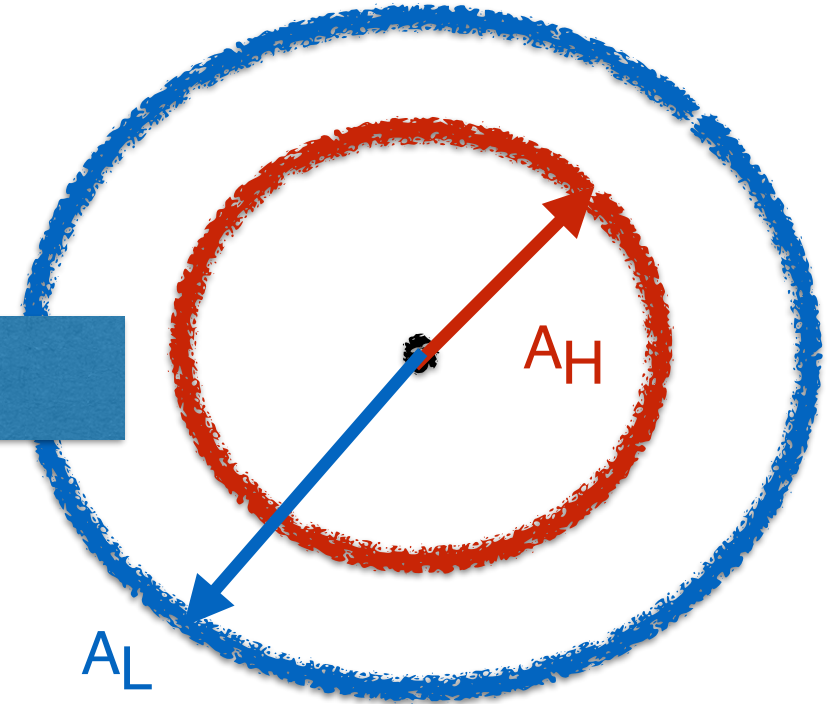




## 7. Fission Yields : systematic uncertainties and errors

$$Y(Z, A) = \frac{\partial N_f}{\partial Z \partial A} \frac{1}{\xi(Z, A, E)} \int dE \int d\theta$$

Detection system



full phase space measurement :

- correction for angular anisotropy
- correction for q-state distribution

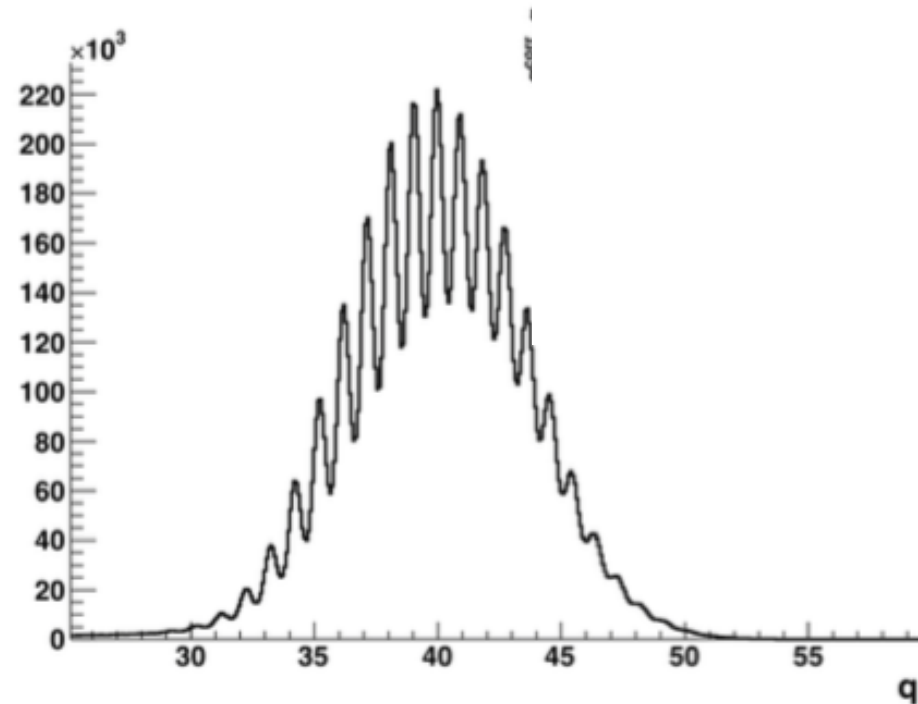
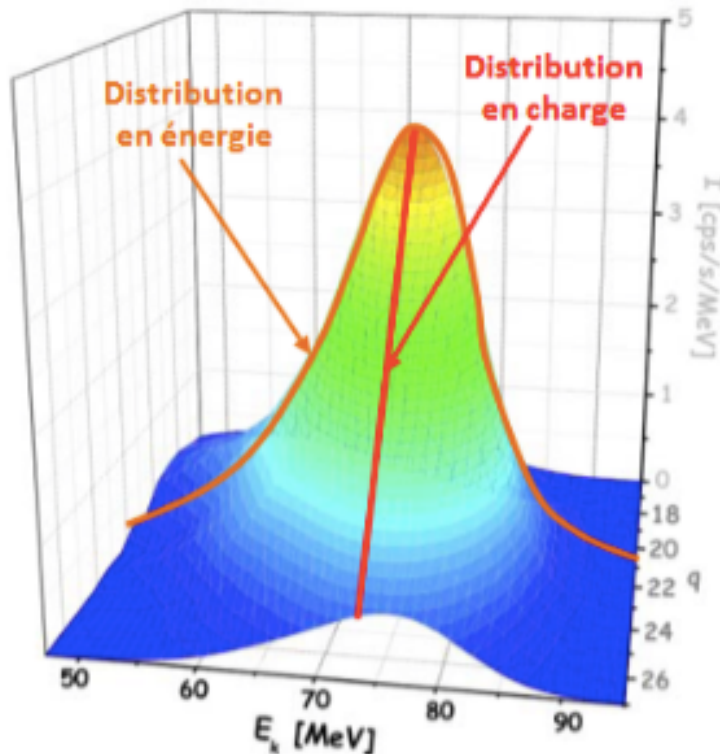
## 7. Fission Yields : systematic uncertainties and errors

In case of magnetic spectrometer : correction for q-state distribution

$$Y(Z, A) = \sum_q \frac{\partial N_f}{\partial Z \partial A}(q) \frac{1}{\xi(Z, A, E, q)} \int dE \int d\theta$$

Lohengrin : estimated

VAMOS : (large acceptance)  
measured





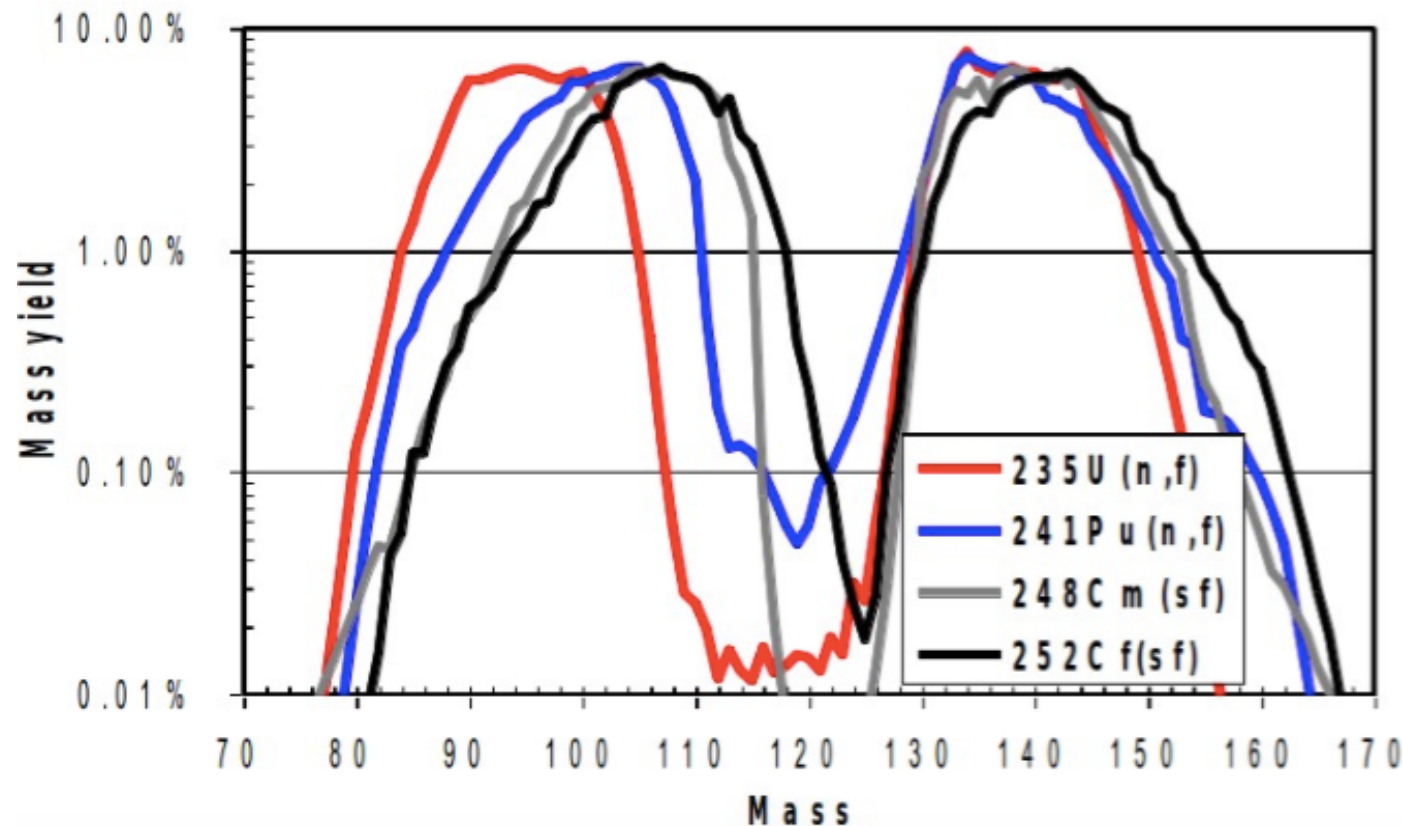
## 7. Fission Yields : systematic uncertainties and errors

Target composition :

Typical actinide target contains several other actinides

$^{235}\text{U}$ : 93.27% ( $^{234}\text{U}$ : 1.05%,  $^{238}\text{U}$ : 5.68%)

$^{239}\text{Pu}$ : 98.41 (40)%,  $^{240}\text{Pu}$ : 1.58(40)%



Need a careful analysis !!

## 8. Conclusions

---

Measurement of fission yields is challenging

Standard techniques based on 2E-2v measurements pursue challenging programmes in mass distributions and neutron multiplicities

gamma spectroscopy is a powerful tool for isotopic identification and fission pair identification

New innovative techniques based on nuclear-induced fission using spectrometers allow for new-generation data on a wide range (and new range) of actinides

This ensemble of data is challenging and constraining for the development of nuclear fission models