



Post – Fukushima Characterization of melted fuel

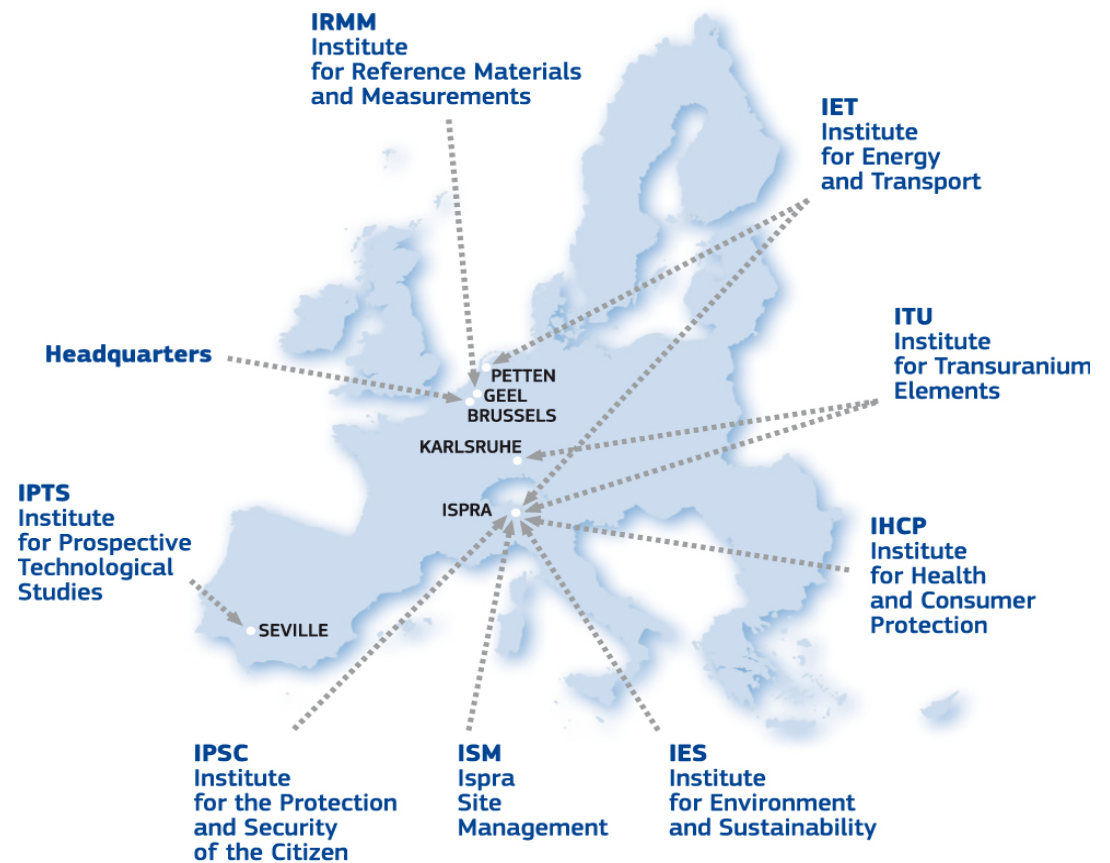
*University of Bologna
5 April 2013*

*P. Schillebeeckx
EC – JRC – IRMM*

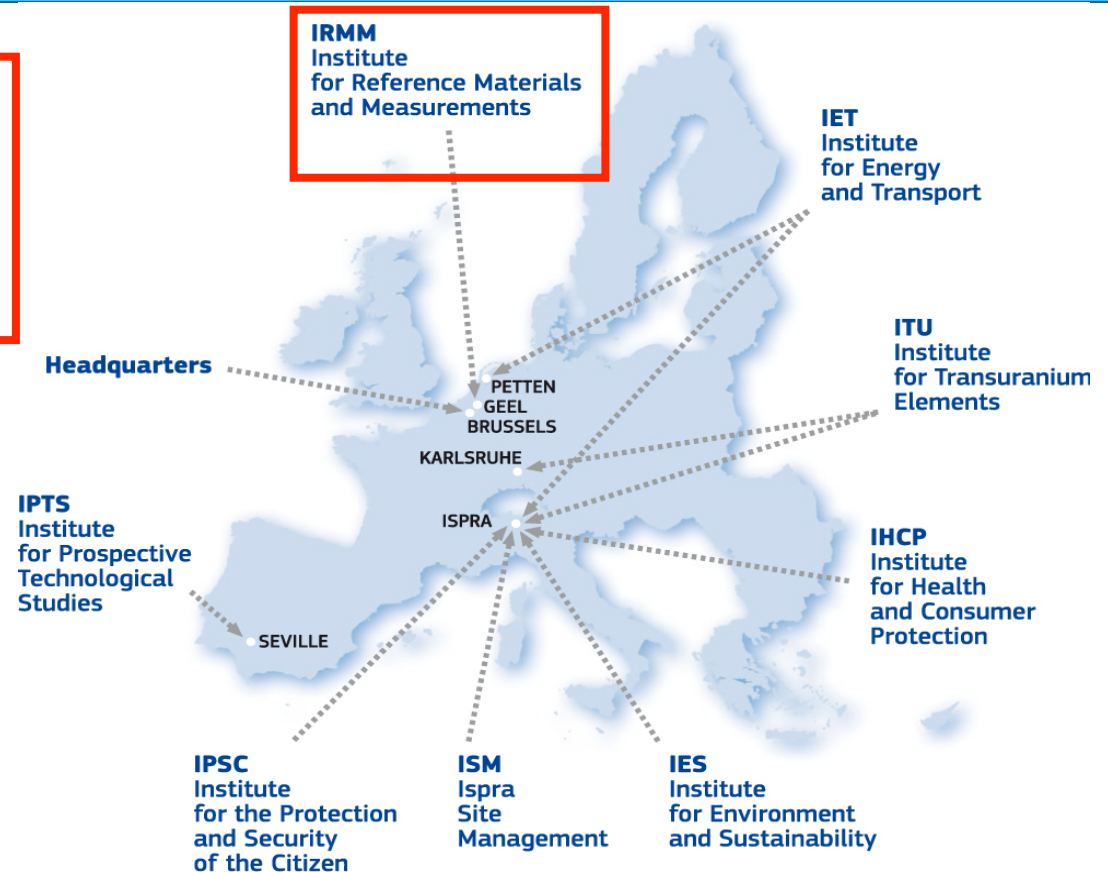
Standards for Nuclear Safety, Security and Safeguards (SN3S)

EC – JRC – IRMM

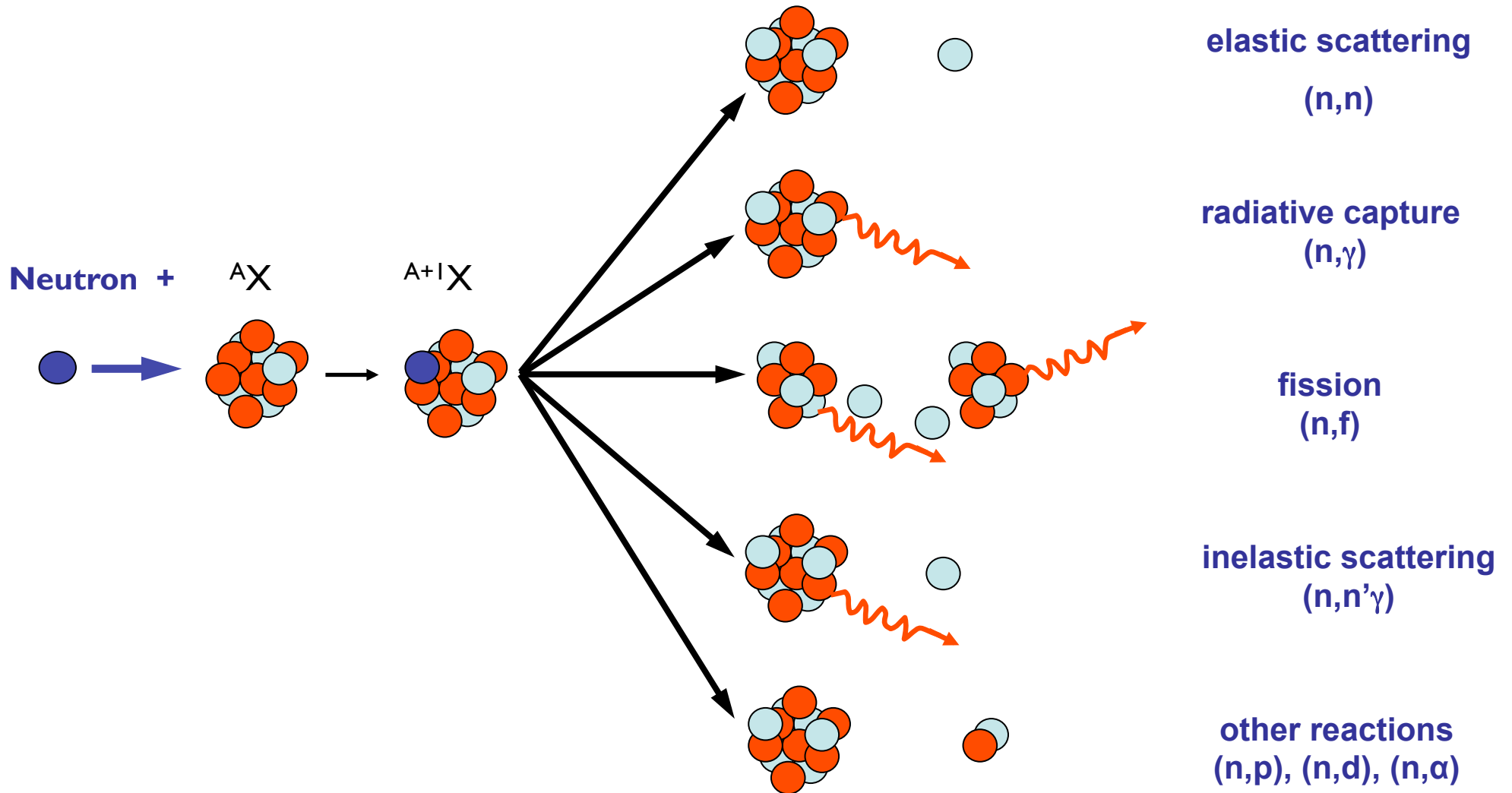
Institute for Reference Materials & Measurements



Mission:
to promote a **common and reliable**
European measurement system
in support of EU policies

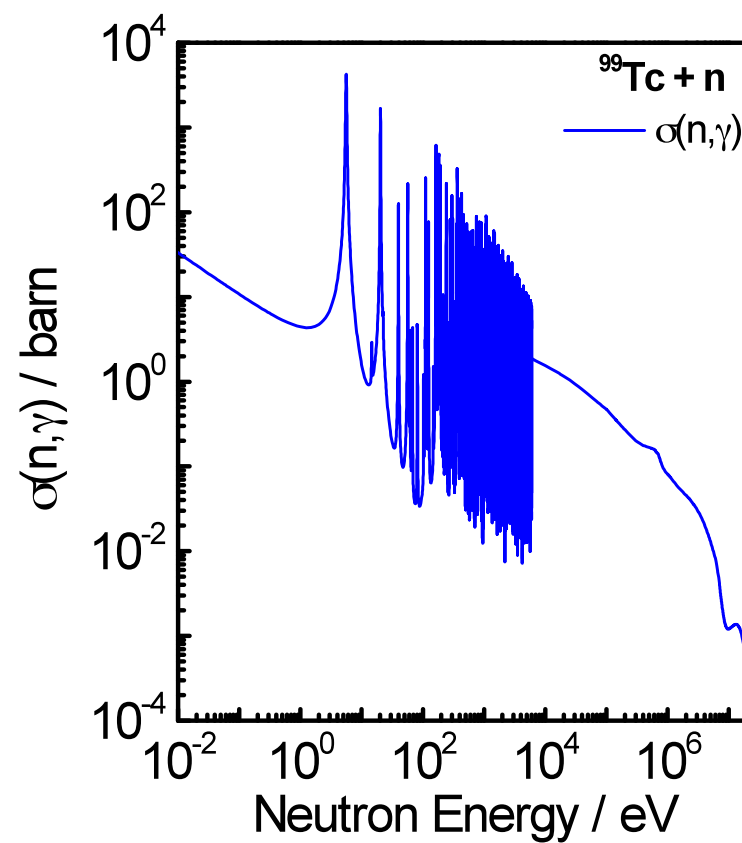


Neutron induced reactions

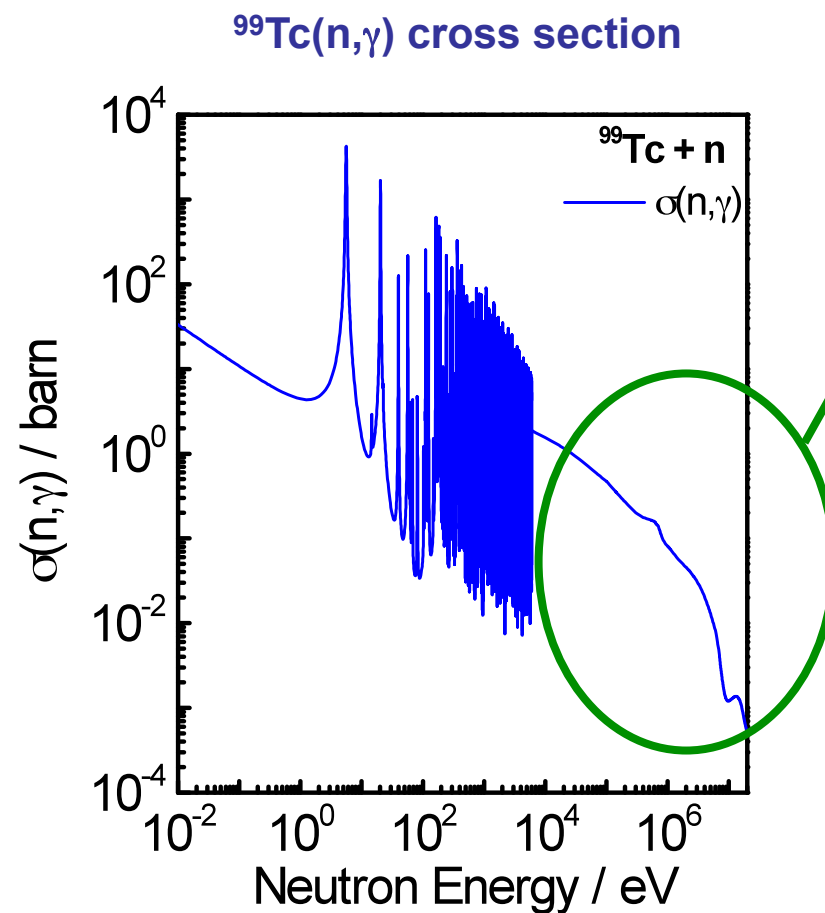


Experimental facilities at IRMM

$^{99}\text{Tc}(n,\gamma)$ cross section



Experimental facilities at IRMM



Van de Graaff



Mono-energetic neutrons
(cp,n) reactions

Experimental facilities at IRMM

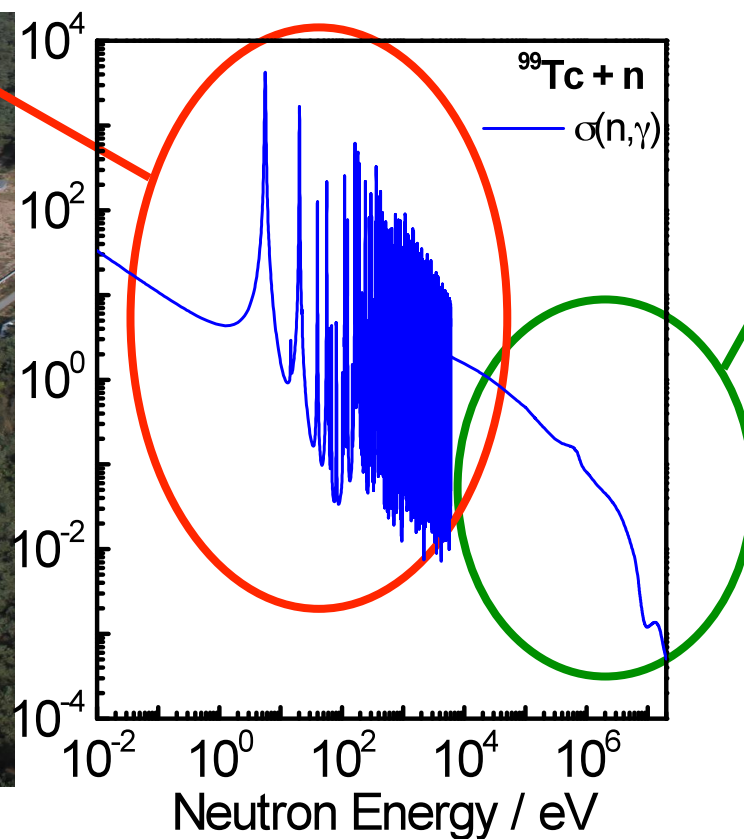


GELINA



**White neutron source
+
Time-of-flight (TOF)**

$^{99}\text{Tc}(n,\gamma)$ cross section



Van de Graaff



**Mono-energetic neutrons
(cp,n) reactions**

TOF - Facility GELINA

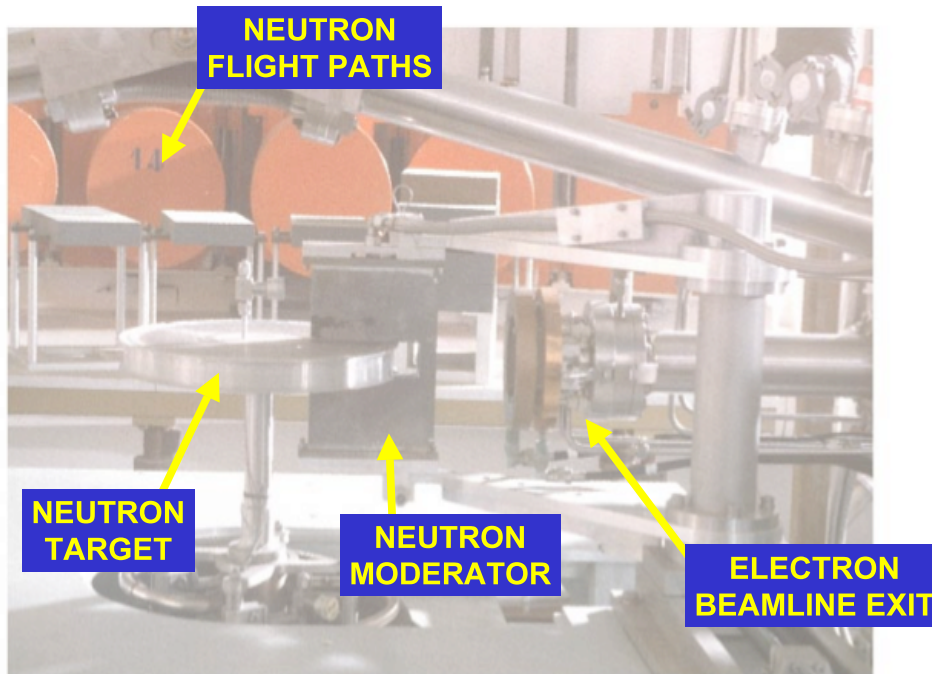


TOF - Facility GELINA



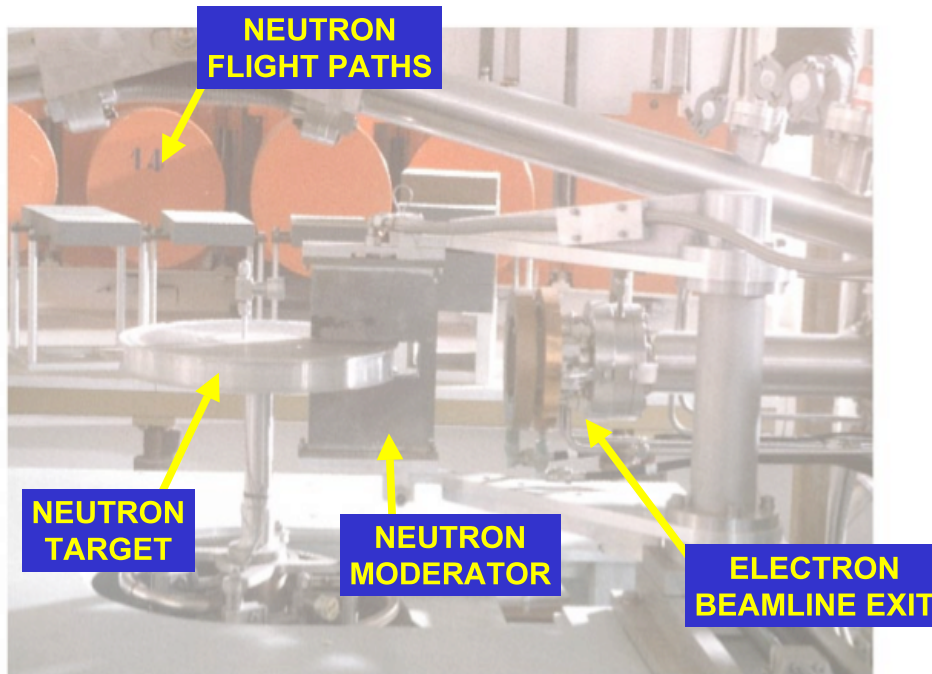
- Pulsed white neutron source
($10 \text{ meV} < E_n < 20 \text{ MeV}$)
- Neutron energy : time-of-flight (TOF)
- Multi-user facility: 10 flight paths
(10 m - 400 m)
- Measurement stations with special equipment to perform:
 - Total cross section measurements
 - Partial cross section measurements

Neutron production

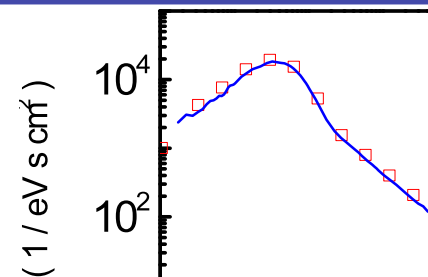


- e^- accelerated to $E_{e-,max} \approx 140$ MeV
- Bremsstrahlung in U-target
(rotating & cooled with liquid Hg)
- (γ, n) , (γ, f) in U-target
- Low energy neutrons by moderation
(water moderator in Be-canning)

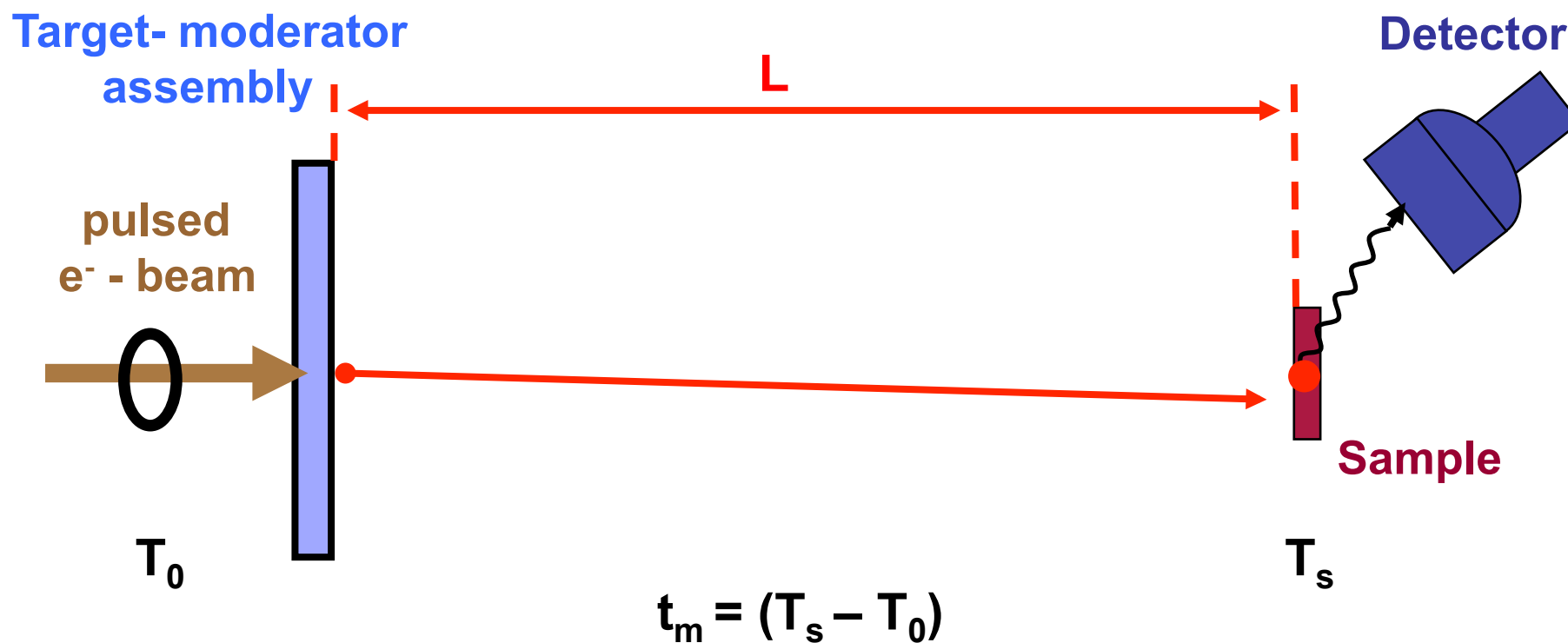
Neutron production



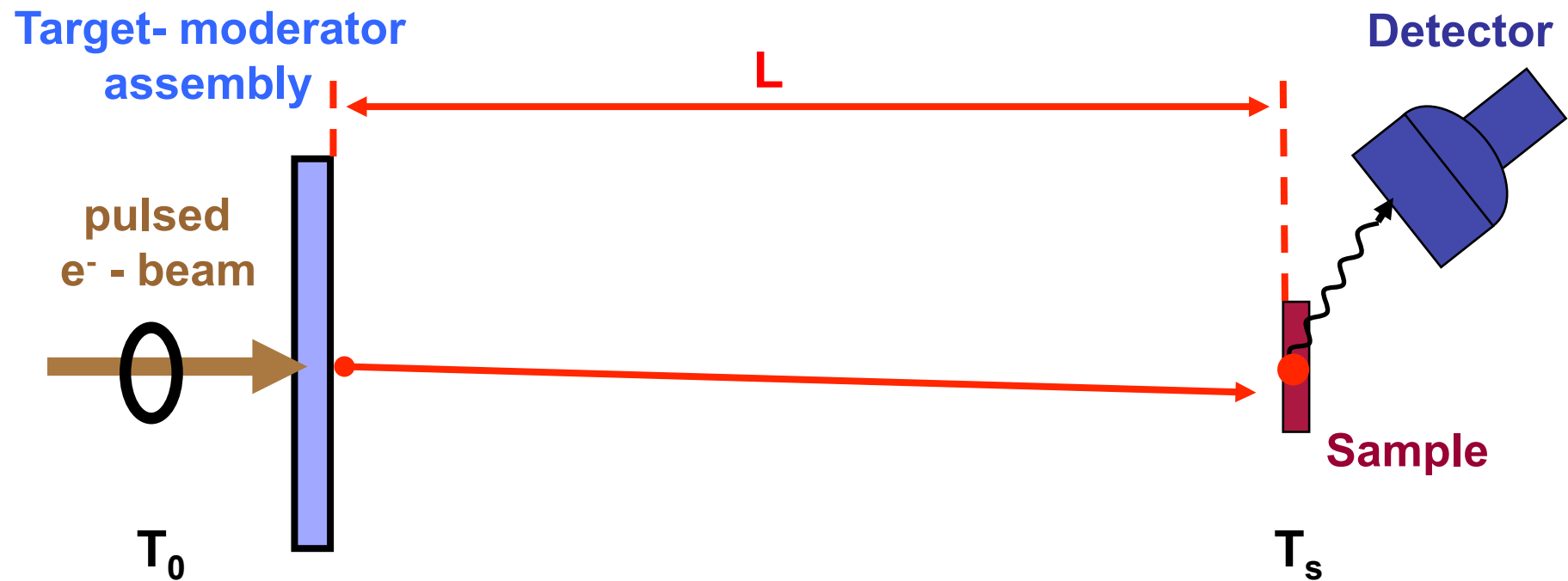
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Time – of – flight technique



Time – of – flight technique

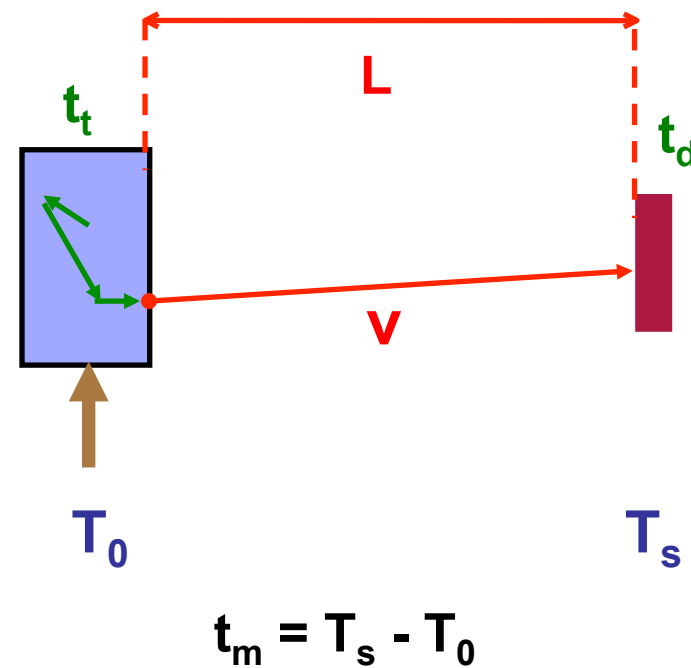


$$t_m = (T_s - T_0)$$

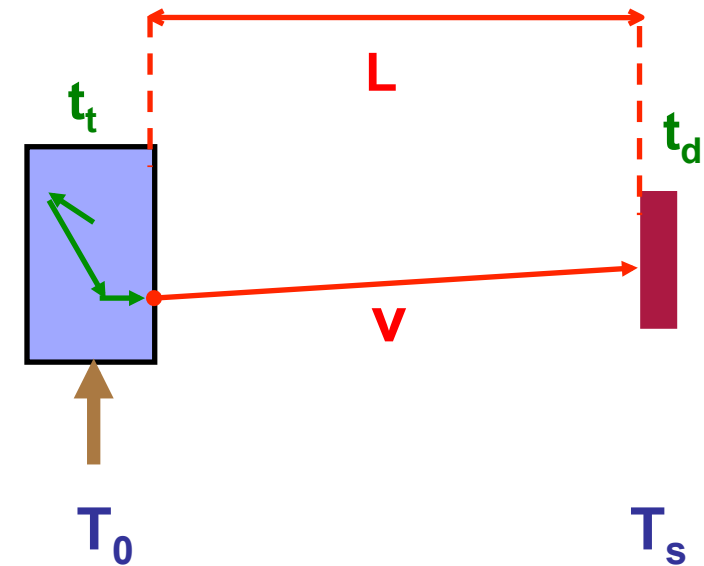
$$v = \frac{L}{t_m}$$

$$E = mc^2 \left(\frac{1}{\sqrt{1 - (v/c)^2}} - 1 \right)$$

Response of TOF-spectrometer



Response of TOF-spectrometer



$$t_m = T_s - T_0$$

$$t = t_m - (t_t + t_d)$$

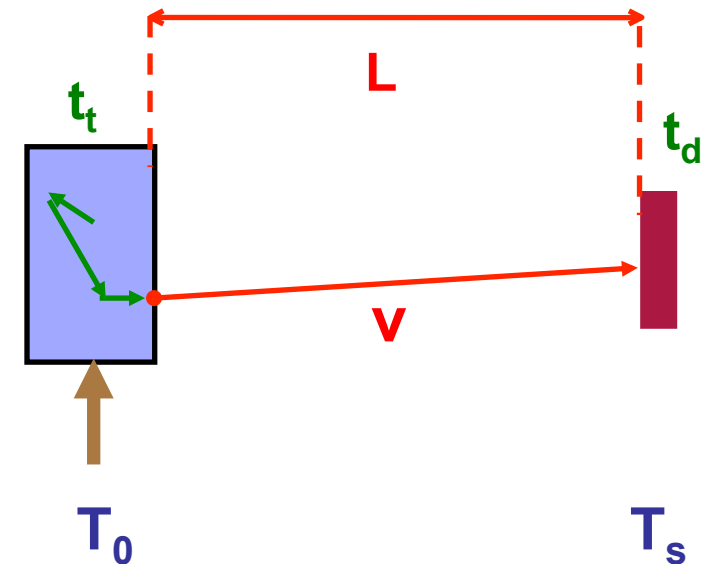
$$v = \frac{L}{t}$$

Response of TOF-spectrometer



$$\frac{\Delta v}{v} = \sqrt{\frac{\Delta t^2}{t^2} + \frac{\Delta L^2}{L^2}} \Rightarrow \frac{\Delta E}{E} = (\gamma + 1)\gamma \frac{\Delta v}{v}$$

- ΔL (~ 1 mm)
- Δt



$$t_m = T_s - T_0$$

$$t = t_m - (t_t + t_d)$$

$$v = \frac{L}{t}$$

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▪ ΔL (~ 1 mm)

▪ Δt

• Initial burst width

ΔT_0

• Time jitter detector & electronics

ΔT_s

• Neutron transport in target - moderator

Δt_t

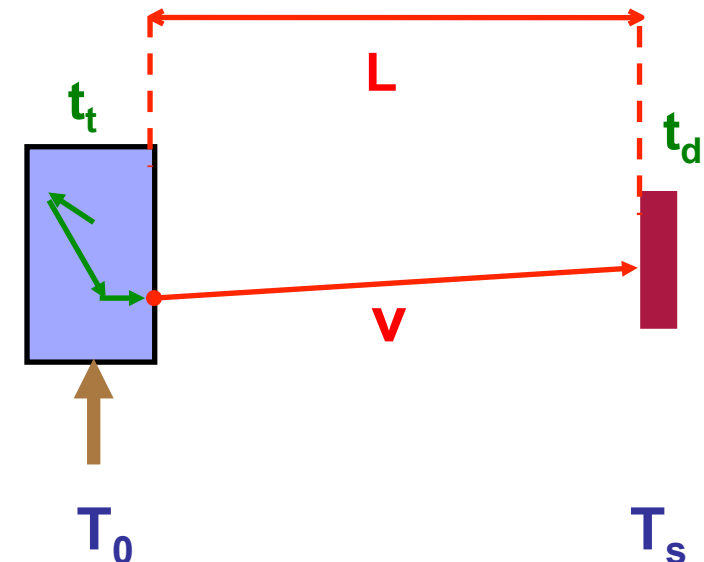
• Neutron transport in detector

Δt_d

$$t_m = T_s - T_0$$

$$t = t_m - (t_t + t_d)$$

$$v = \frac{L}{t}$$



Response of TOF-spectrometer

$$\frac{\Delta v}{v} = \sqrt{\frac{\Delta t^2}{t^2} + \frac{\Delta L^2}{L^2}} \Rightarrow \frac{\Delta E}{E} = (\gamma + 1)\gamma \frac{\Delta v}{v}$$

▪ ΔL (~ 1 mm)

▪ Δt

• Initial burst width

ΔT_0

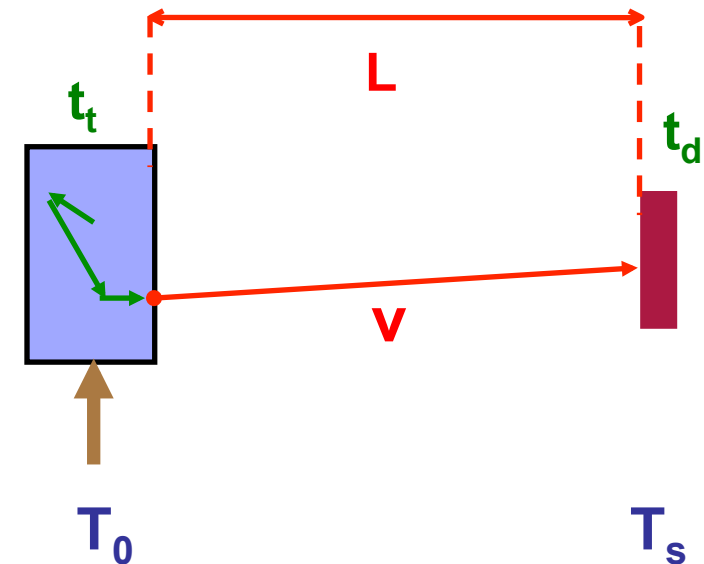
• Time jitter detector & electronics

ΔT_s

• Neutron transport in target - moderator Δt_t

• Neutron transport in detector

Δt_d

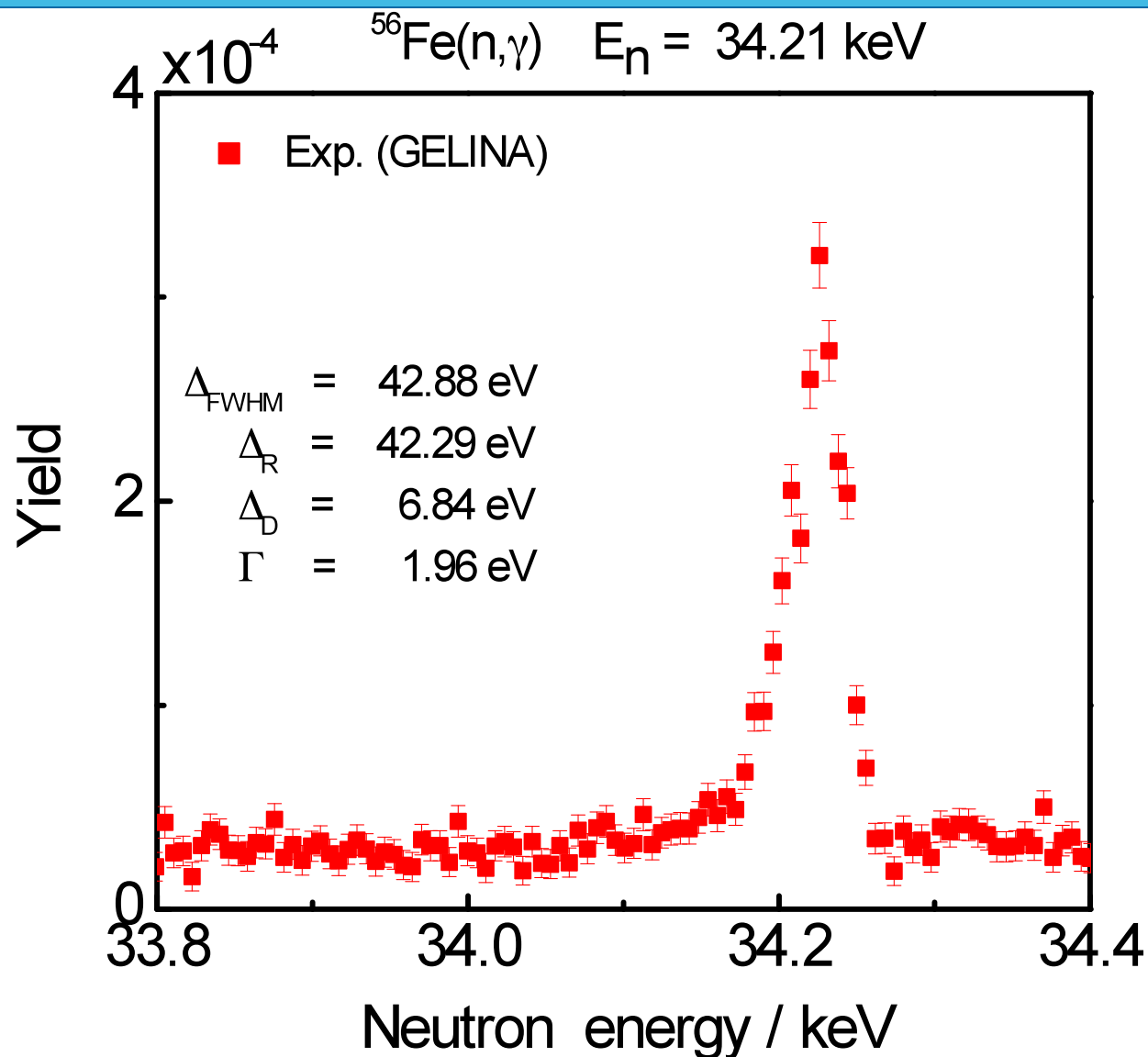


$$t_m = T_s - T_0$$

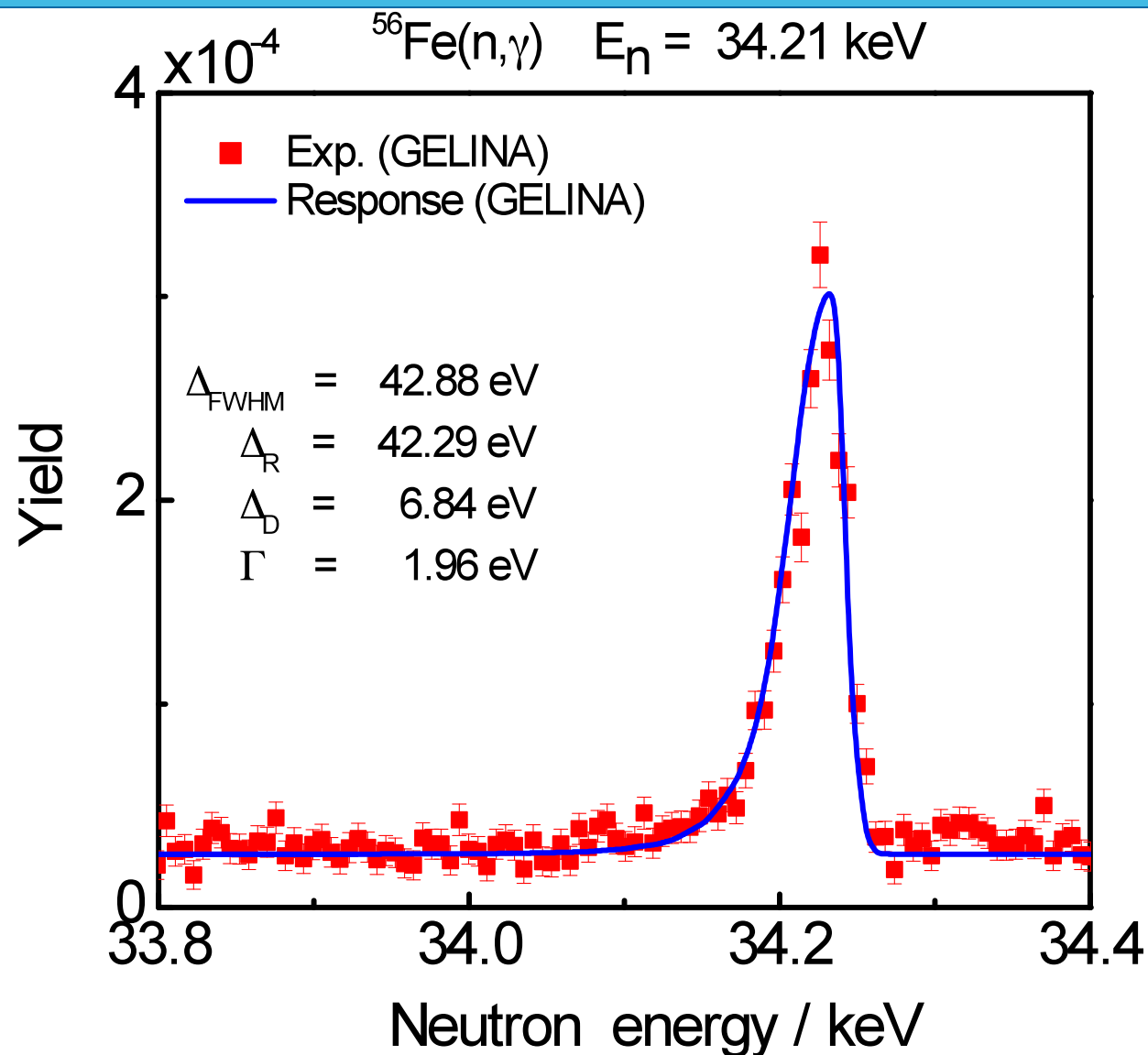
$$t = t_m - (t_t + t_d)$$

$$v = \frac{L}{t}$$

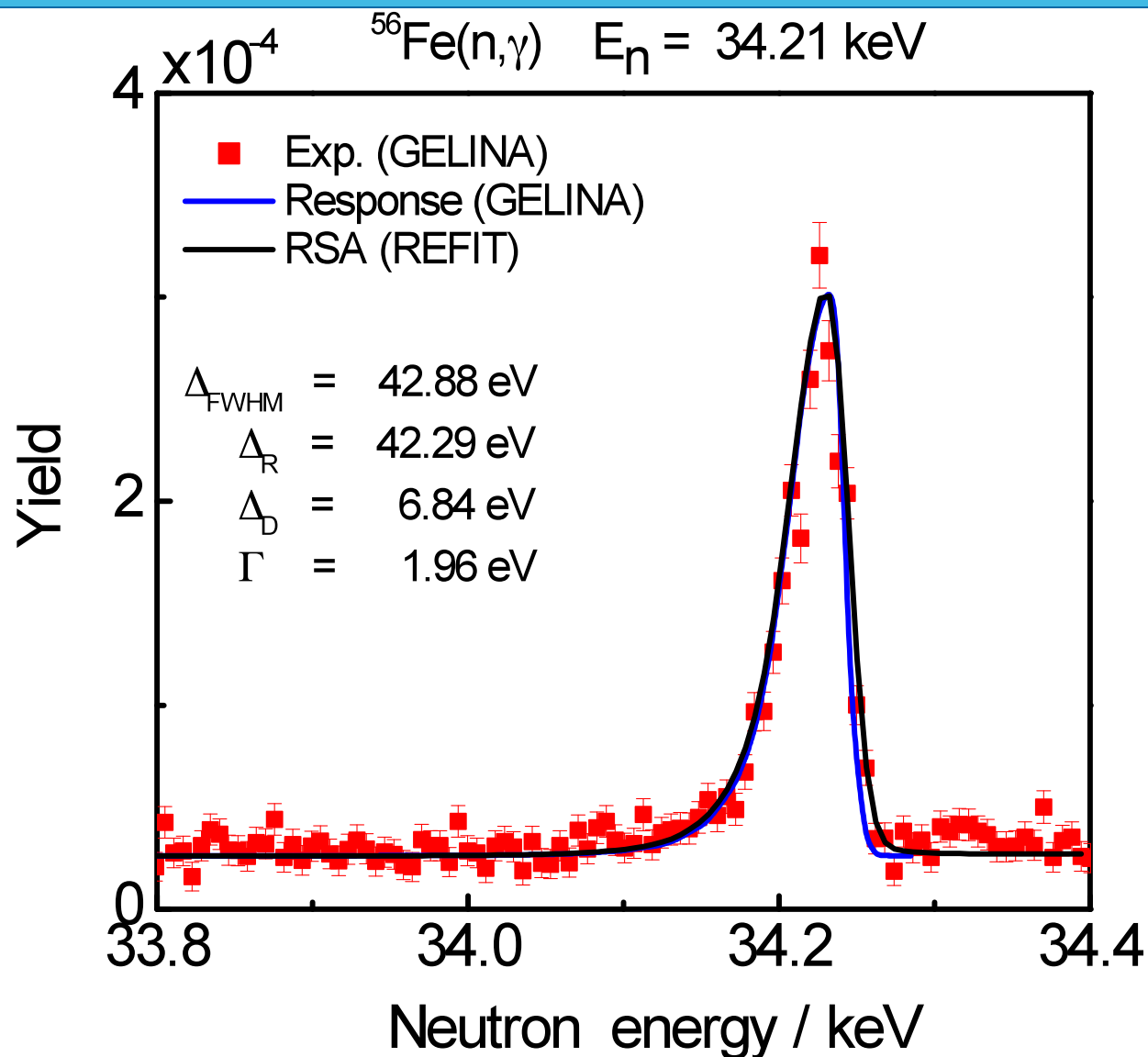
Response GELINA (60 m)



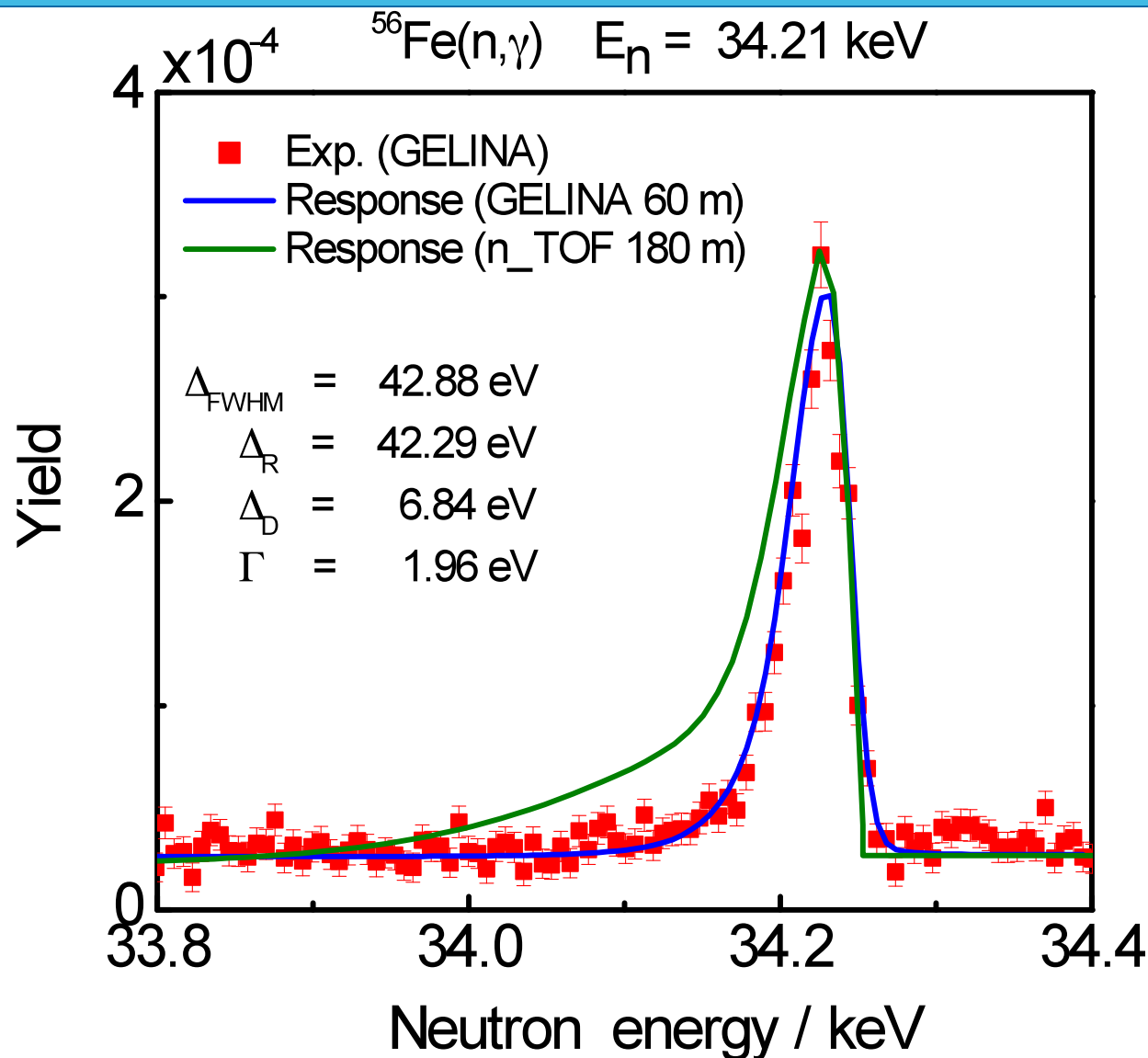
Response GELINA (60 m)



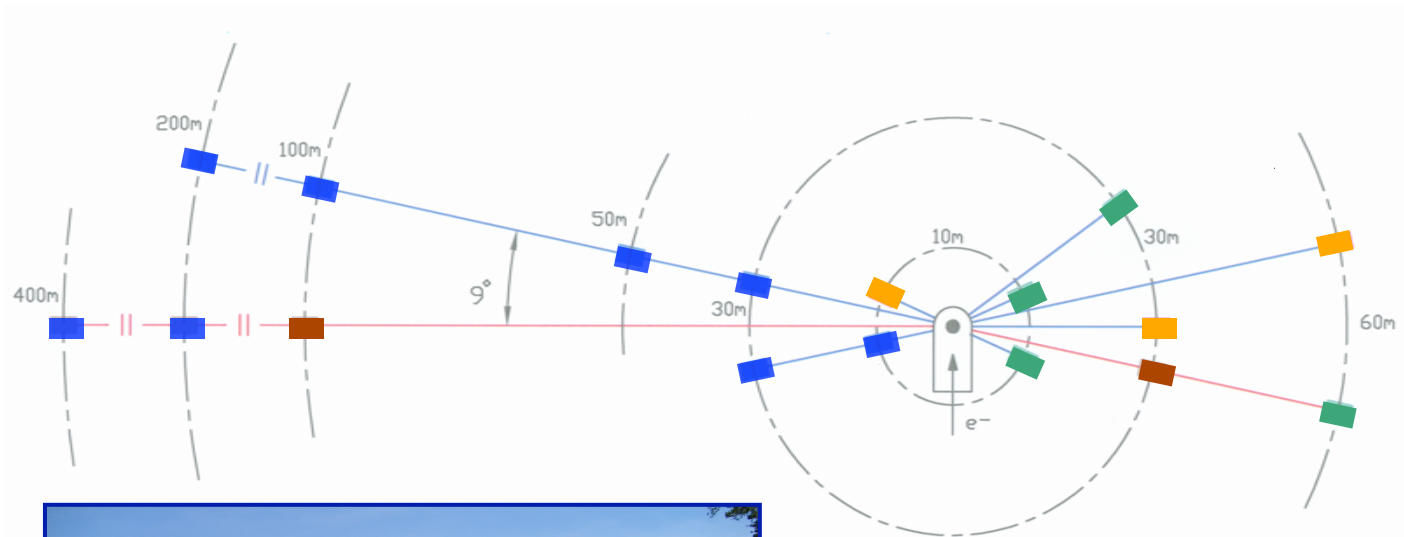
Response GELINA (60 m)



Response GELINA (60 m) < - > n_TOF (180 m)

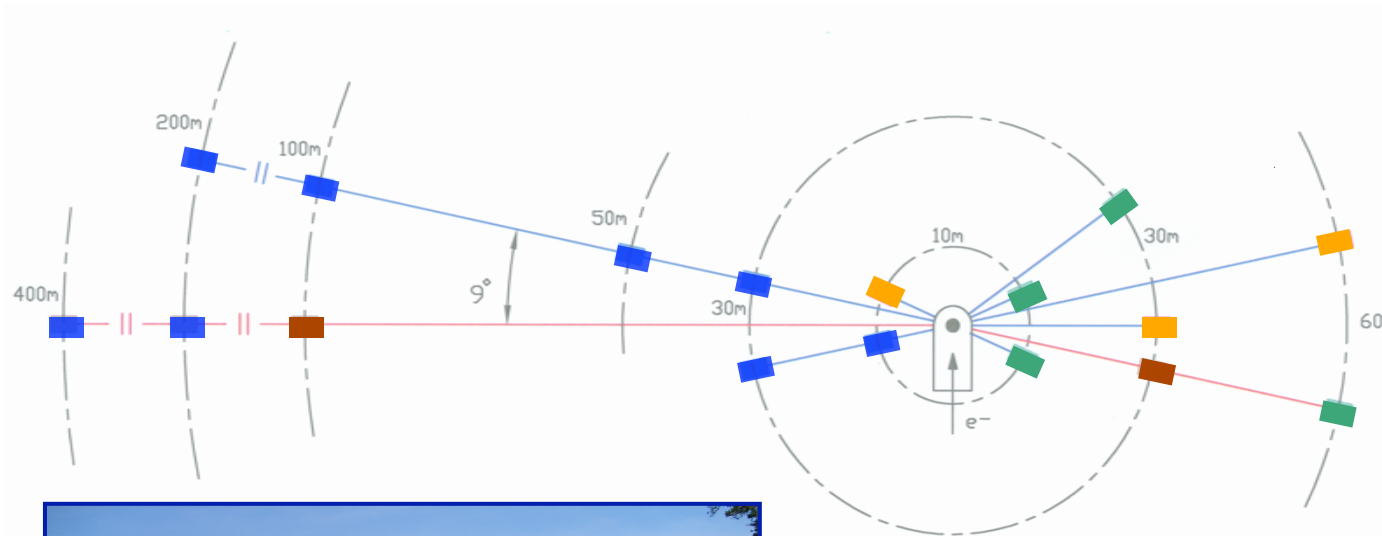


TOF - measurements



- (n,γ)
- (n,tot)
- (n,f) and (n,cp)
- $(n,n'\gamma)$

TOF - measurements



- (n,γ)
- (n,tot)
- (n,f) and (n,cp)
- (n,n'γ)

▪ Velocity from TOF

$$v = \frac{L}{t}$$

▪ Neutron flux $\Rightarrow L \searrow$

$$\varphi(L) \propto \frac{1}{L^2}$$

▪ Resolution $\Rightarrow L \nearrow$

$$\begin{aligned} \frac{\Delta v}{v} &= \sqrt{\frac{\Delta t^2}{t^2} + \frac{\Delta L^2}{L^2}} \\ &= \frac{1}{L} \sqrt{v^2 \Delta t^2 + \Delta L^2} \end{aligned}$$

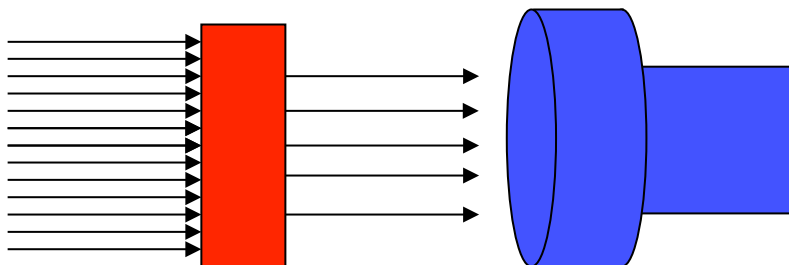
Cross section measurements

Transmission : $\sigma(n, \text{tot})$

$$T \cong e^{-n \sigma_{\text{tot}}}$$

T : transmission

Fraction of the neutron beam traversing the sample without any interaction



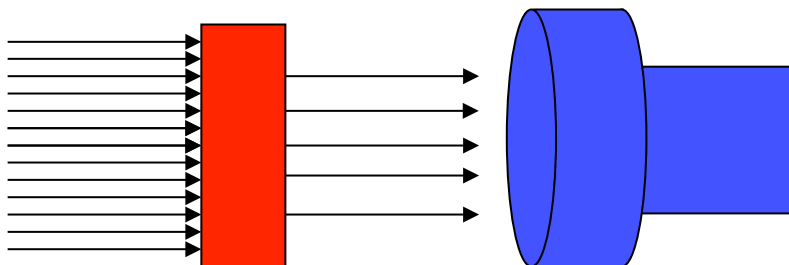
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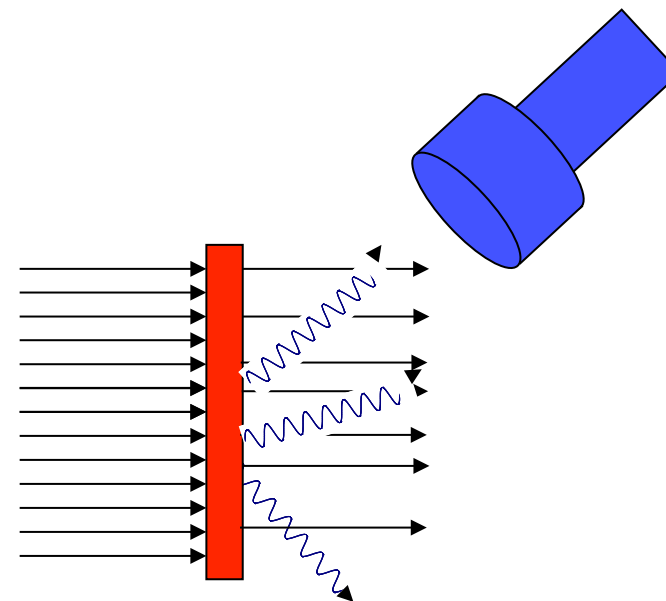


Reaction yield : $\sigma(n, r)$

$$Y_r \approx (1 - e^{-n \sigma_{\text{tot}}}) \frac{\sigma_r}{\sigma_{\text{tot}}}$$

Y_r : reaction yield

Fraction of the neutron beam creating a (n,r) reaction in the sample



Transmission measurements at GELINA



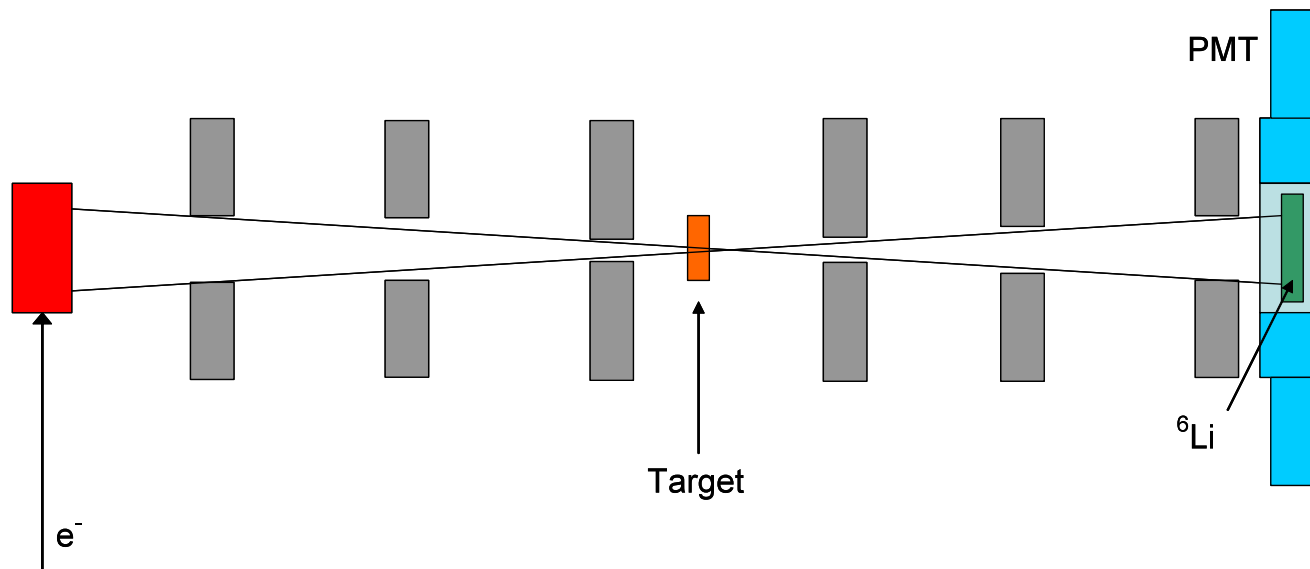
$$T = \frac{C_{in}}{C_{out}} \propto e^{-n \sigma_{tot}}$$

- (1) All detected neutrons passed through the sample**
- (2) Neutrons scattered in the target do not reach detector**
- (3) Sample perpendicular to parallel neutron beam**

Transmission measurements at GELINA

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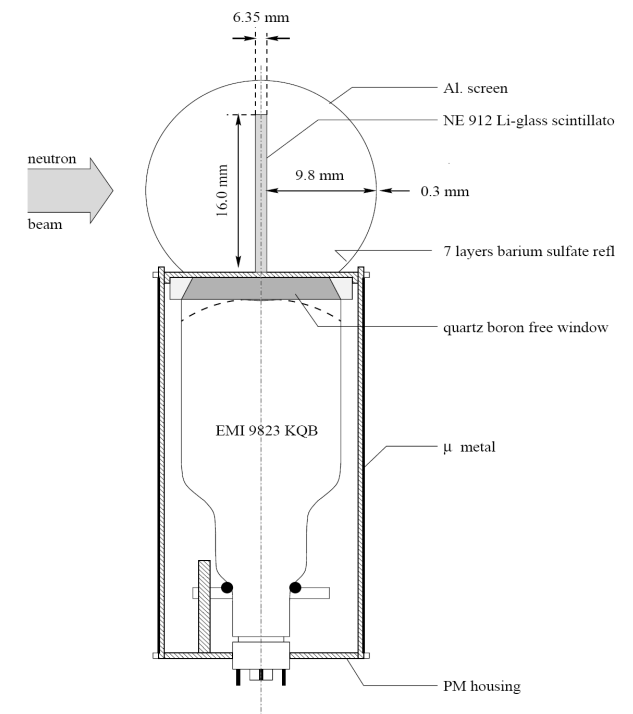
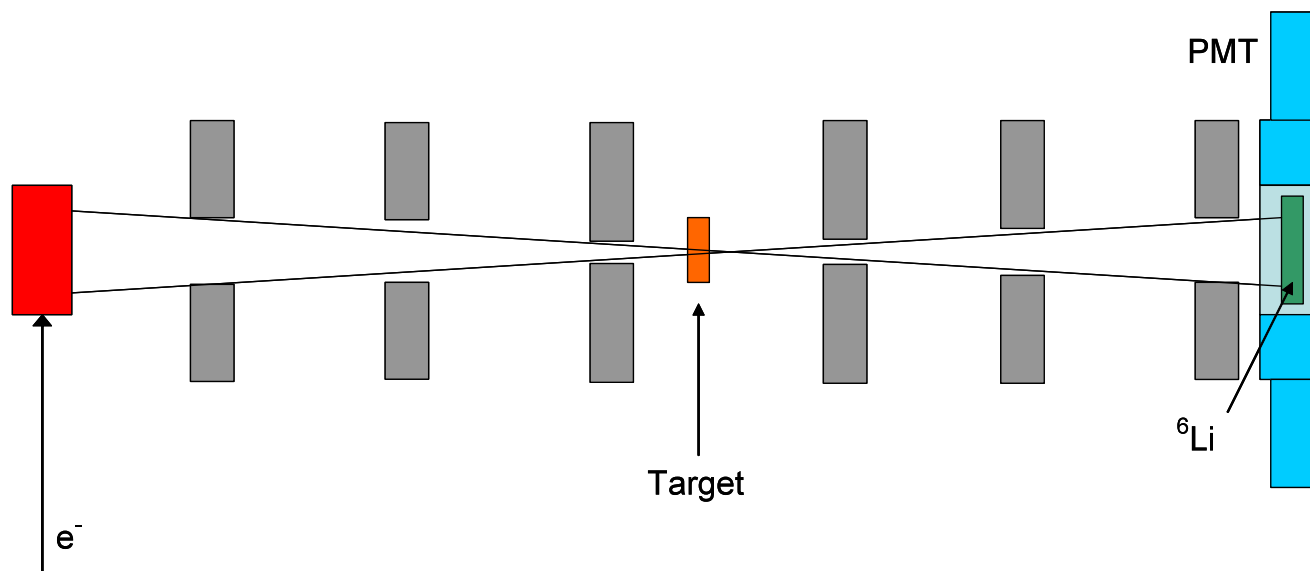
- (1) All detected neutrons passed through the sample
- (2) Neutrons scattered in the target do not reach detector
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 - ⇒ Good transmission geometry (collimation)
 - ⇒ Homogeneous target



Transmission measurements at GELINA

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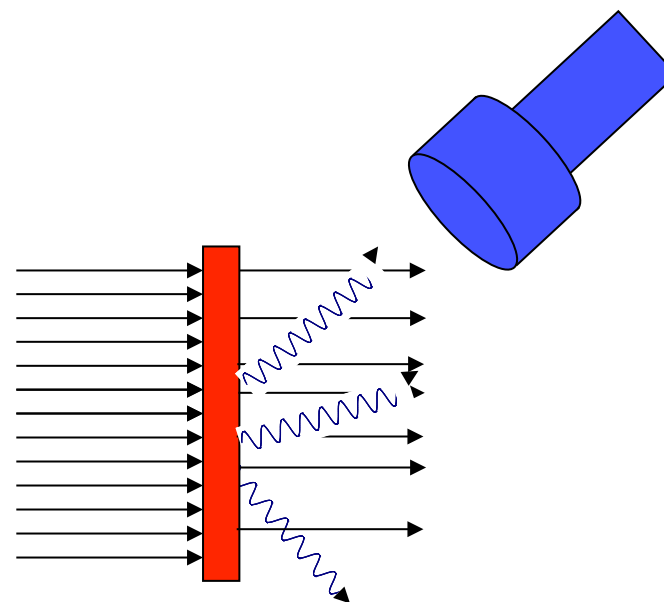


Reaction cross section measurements

Reaction e.g. (n, γ)

$$Y_{\gamma} \approx (1 - e^{-n \sigma_{\text{tot}}}) \frac{\sigma_{\gamma}}{\sigma_{\text{tot}}}$$

$$Y_{\gamma, \text{exp}} = \frac{C_{\gamma}}{\varepsilon_{\gamma}^{\text{eff}} \varphi}$$



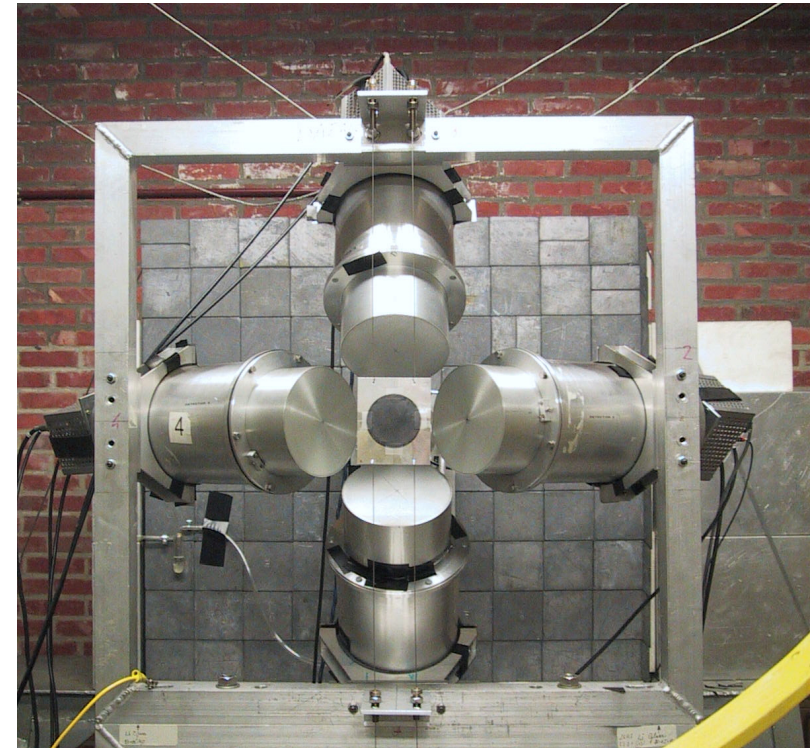
$\sigma(n,\gamma)$ measurements at GELINA



L = 10 m, 30 m and 60 m

Total energy detection principle

- C_6D_6 liquid scintillators
 - 125°
 - PHWT



Collaboration INFN Legnaro (P. Mastinu)
Improvement of C_6D_6 detectors

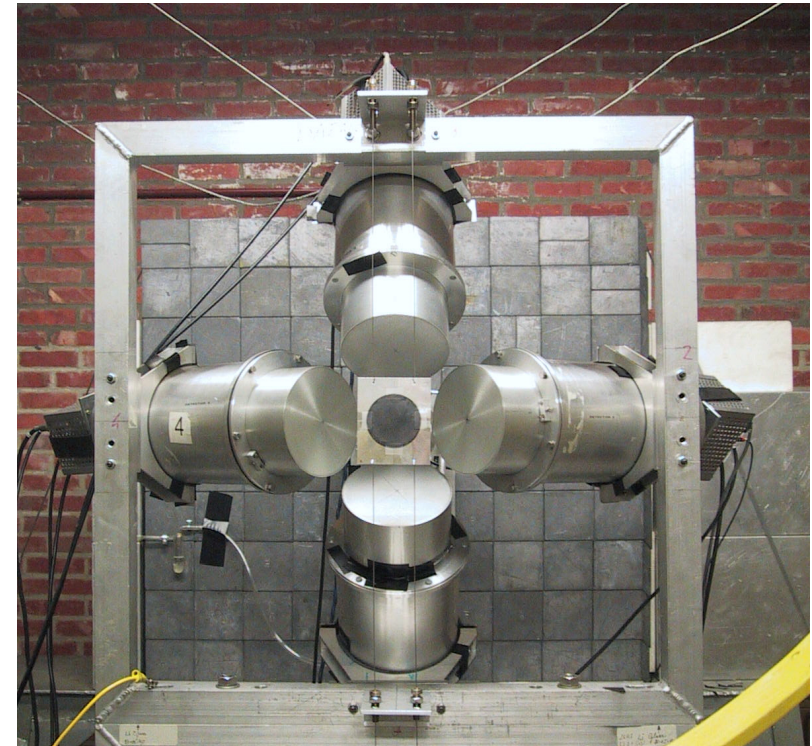
$\sigma(n,\gamma)$ measurements at GELINA

Total energy detection principle

- C_6D_6 liquid scintillators
 - 125°
 - PHWT
- Flux measurements (IC)
 - $^{10}\text{B}(n,\alpha)$
 - $^{235}\text{U}(n,f)$



$L = 10 \text{ m}, 30 \text{ m} \text{ and } 60 \text{ m}$



$\sigma(n,\gamma)$ measurements at GELINA

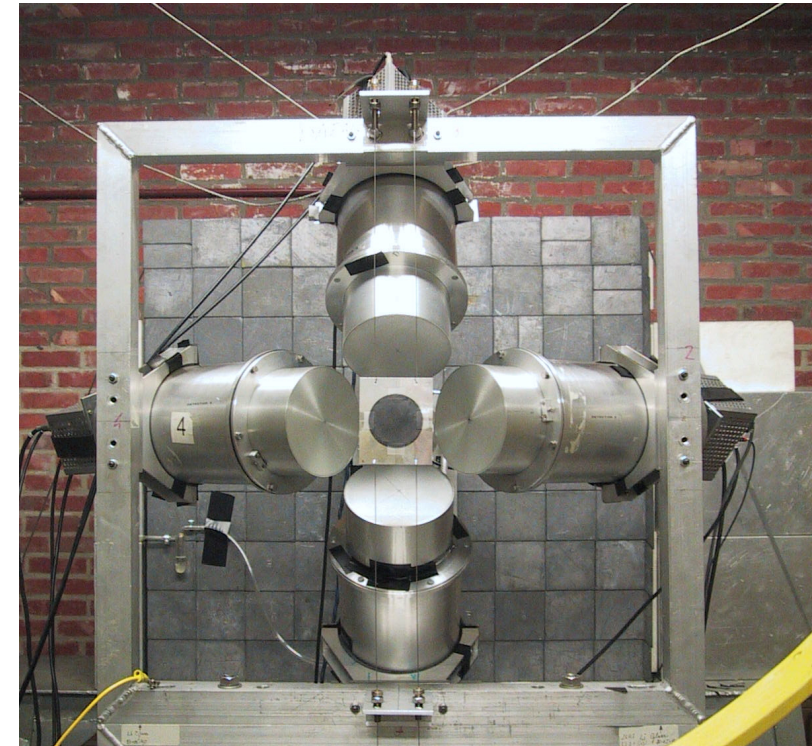
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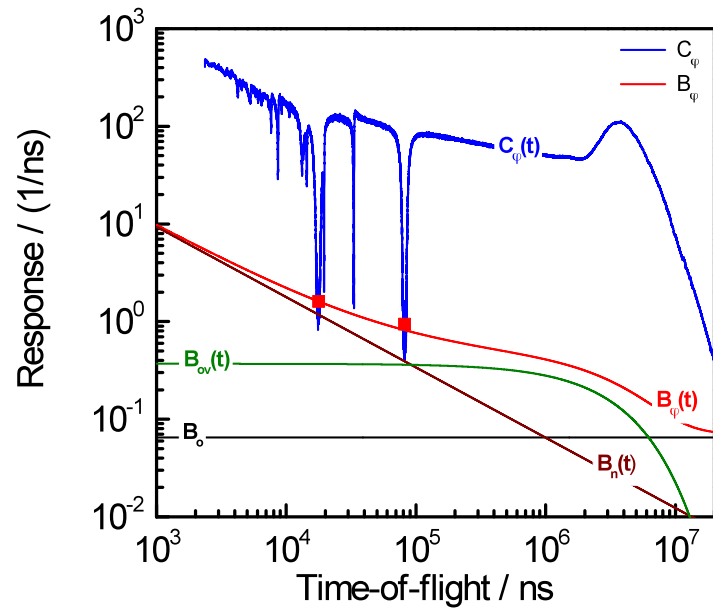


$$Y_{\text{exp}} = N Y_{\varphi} \frac{C_w - B_w}{C_{\varphi} - B_{\varphi}}$$

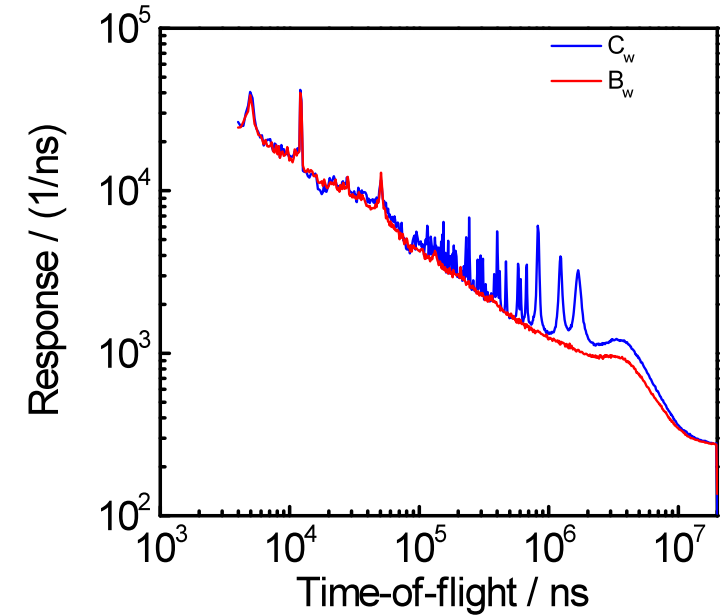
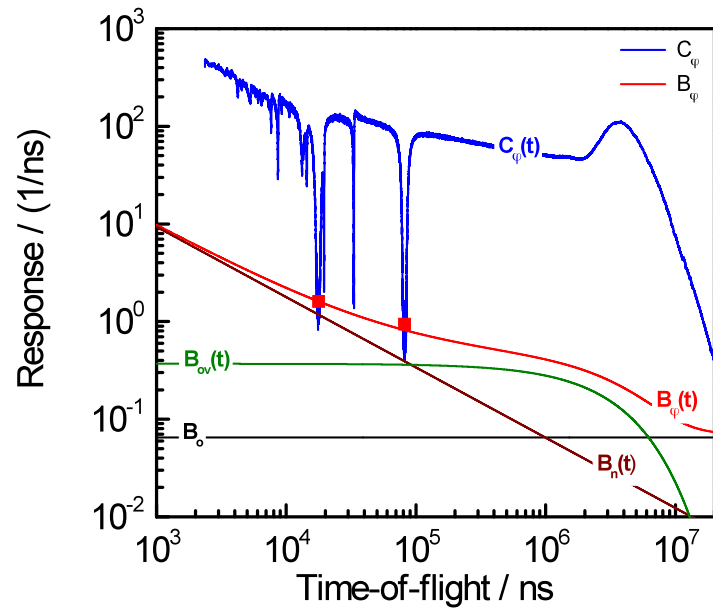
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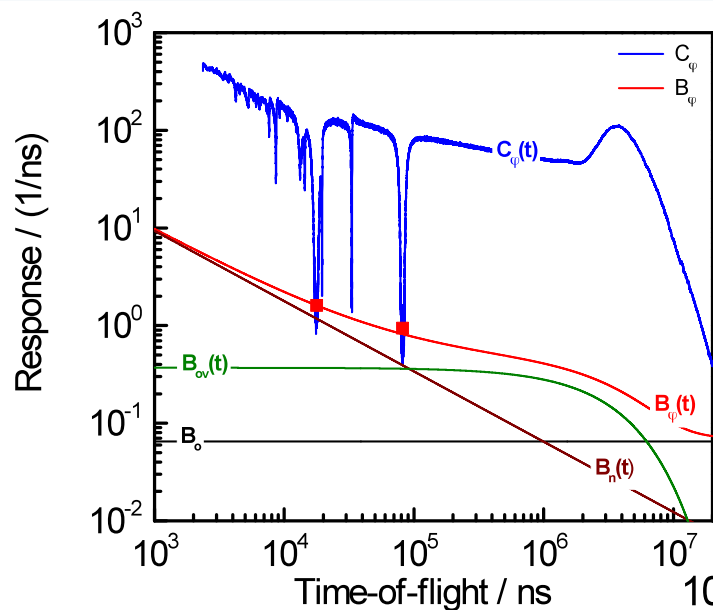
$^{241}\text{Am} + n$ capture at GELINA



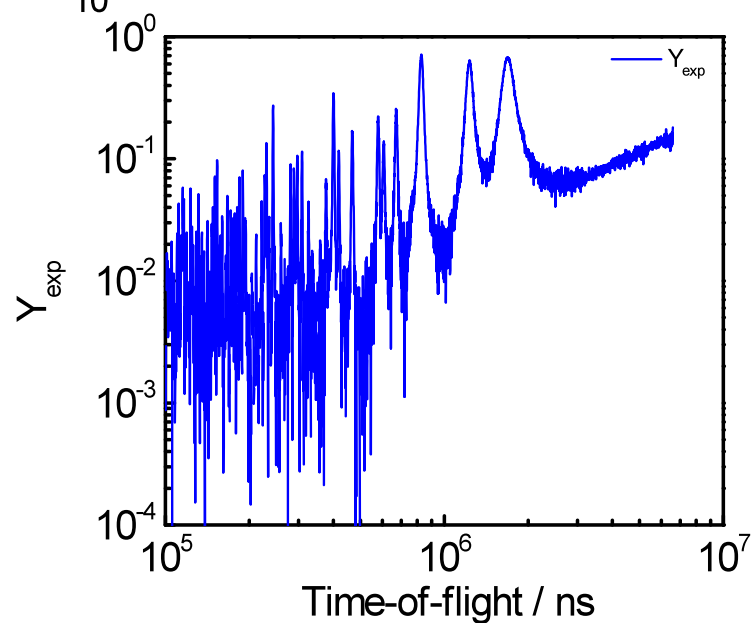
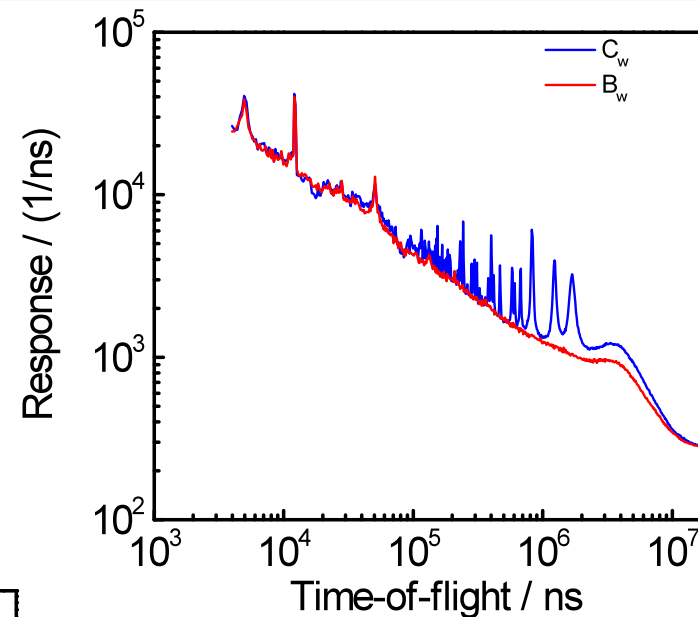
$^{241}\text{Am} + n$ capture at GELINA



$^{241}\text{Am} + n$ capture at GELINA



$$Y_{\text{exp}} = N \frac{C_w - B_w}{C_\varphi - B_\varphi} Y_\varphi$$

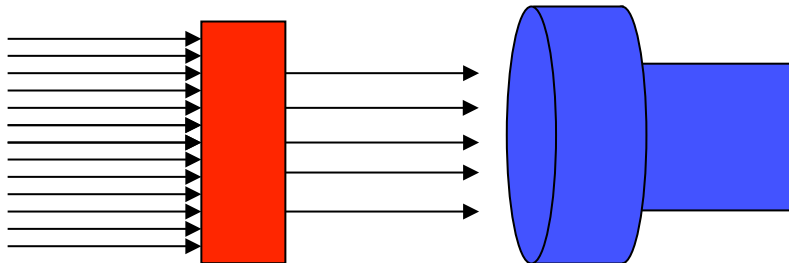


Cross section measurements

Transmission

$$T \cong e^{-n \sigma_{\text{tot}}}$$

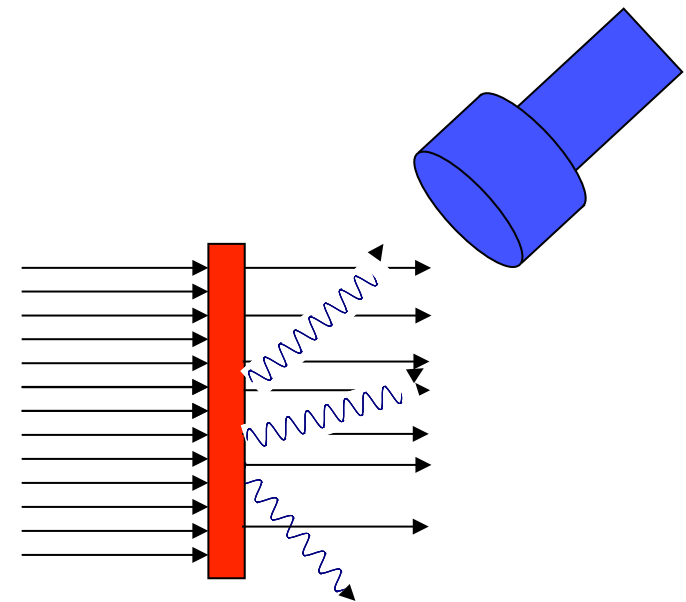
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Fitting model parameters to experimental data



$$\chi^2(\vec{\eta}, \vec{\kappa}) = (\vec{Z}_{\text{exp}} - \vec{Z}_M(\vec{\eta}, \vec{\kappa}))^T \underline{V}_{\vec{Z}_{\text{exp}}}^{-1} (\vec{Z}_{\text{exp}} - \vec{Z}_M(\vec{\eta}, \vec{\kappa}))$$

\vec{Z}_{exp} : experimental observable

$\vec{Z}_M(\vec{\eta}, \vec{\kappa})$: model prediction

$$Z_M(t_m) = \int R(t_m, E_n) Z(E_n) dE_n$$

$\vec{\eta}$ = resonance parameters

- R_j
- J^π
- $(E_R, \Gamma_{n,j}, \Gamma_{\gamma,j}, \Gamma_{f,j}, \dots) j = 1, \dots, n$

$\vec{\kappa}$ = experimental parameters

- Flight Path length
- Resolution
- Sample characteristics
- Doppler (sample temperature)
- Detector characteristics
- ...

Fitting model parameters to experimental data



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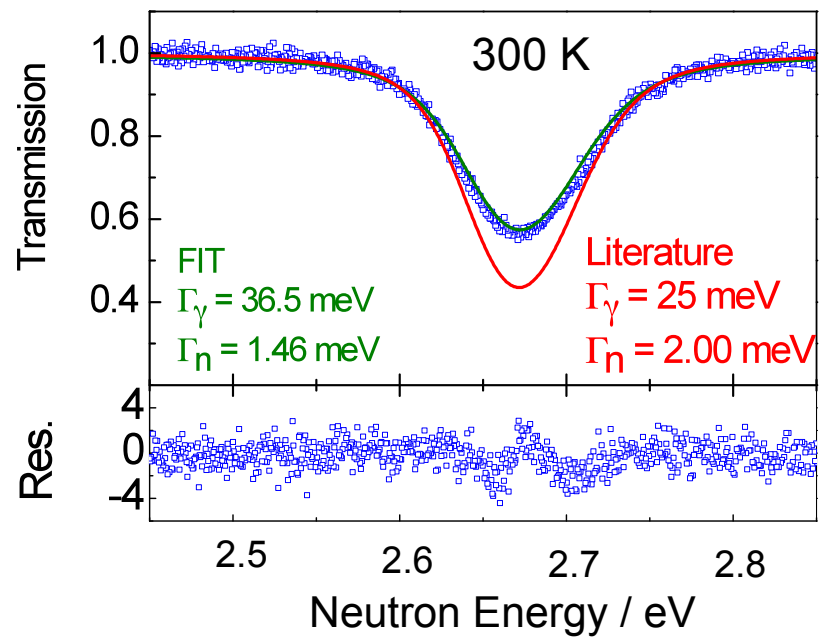
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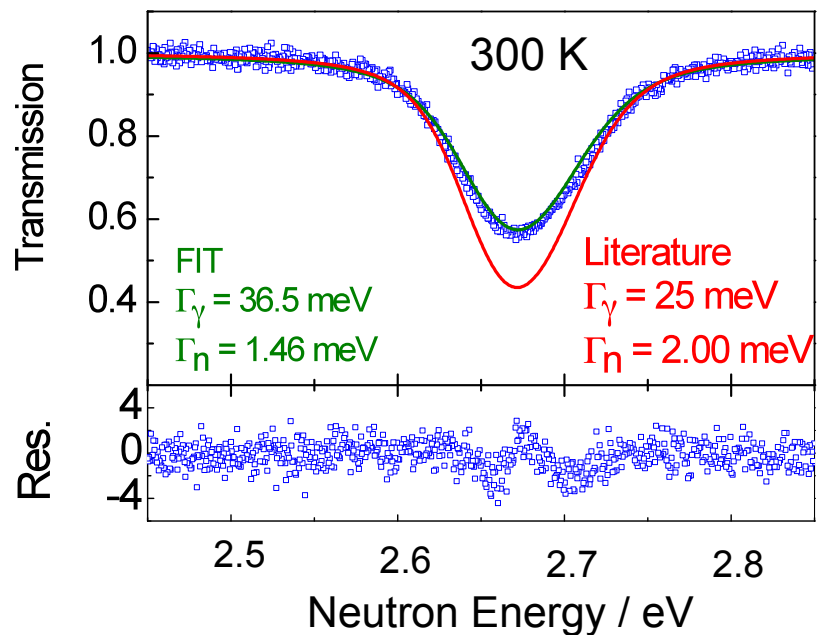
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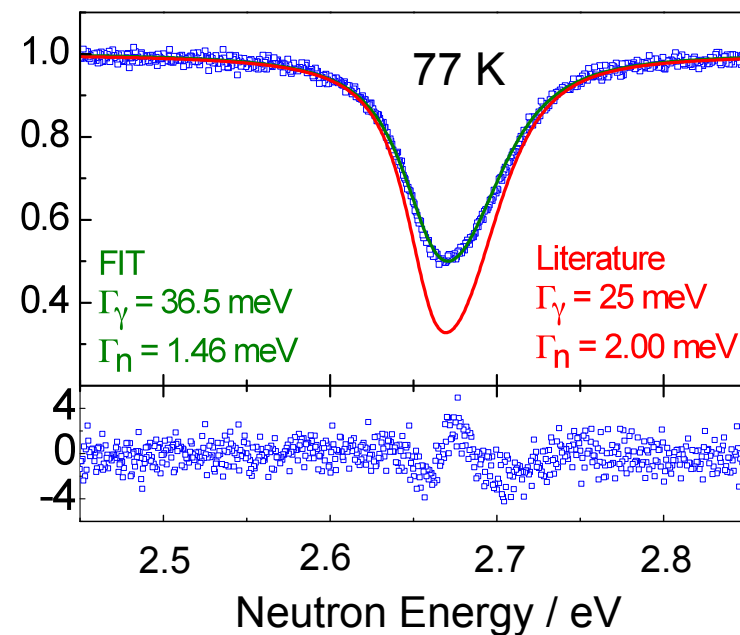
Sample properties: $^{242}\text{PuO}_2$ + carbon powder



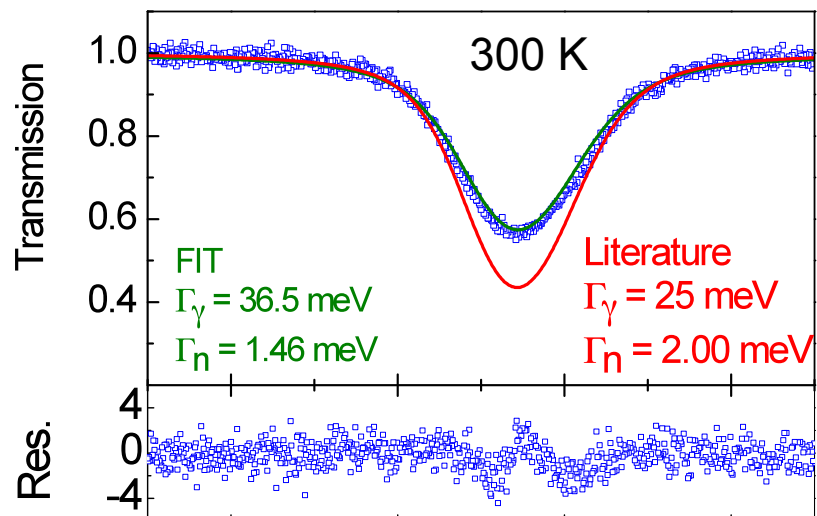
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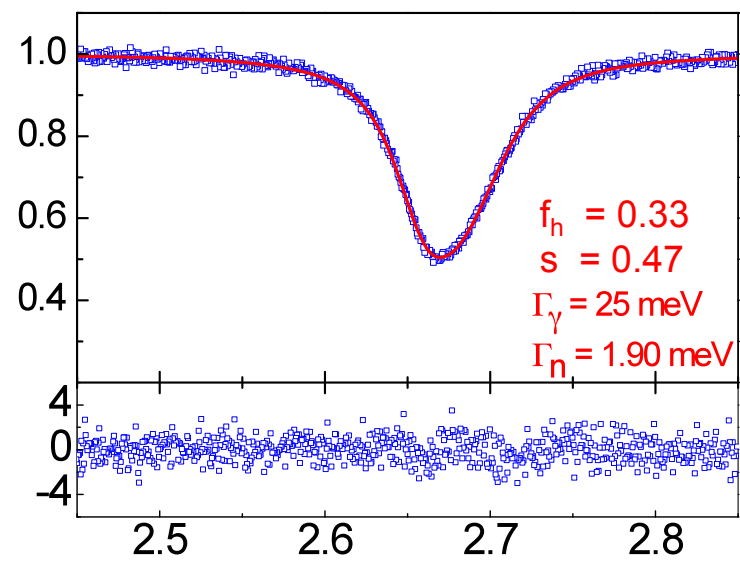
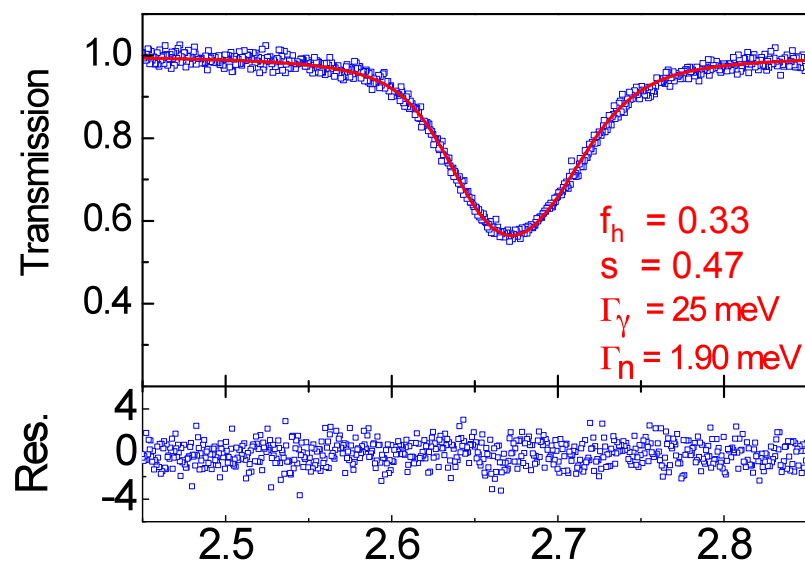
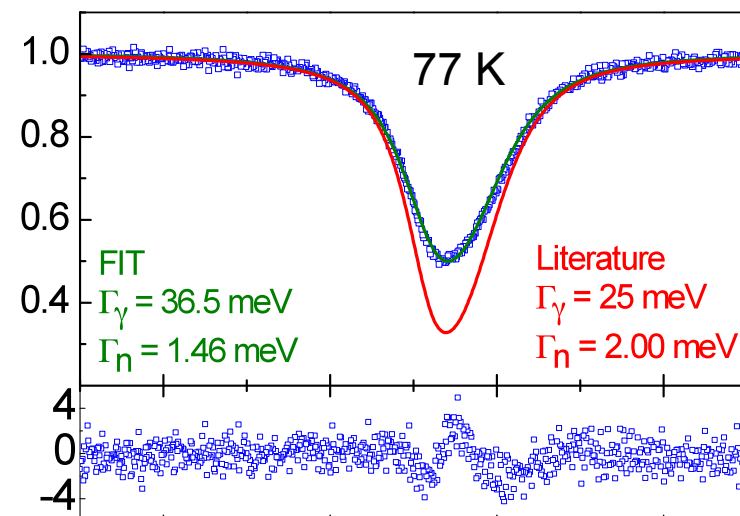
Γ_n underestimated
 Γ overestimated



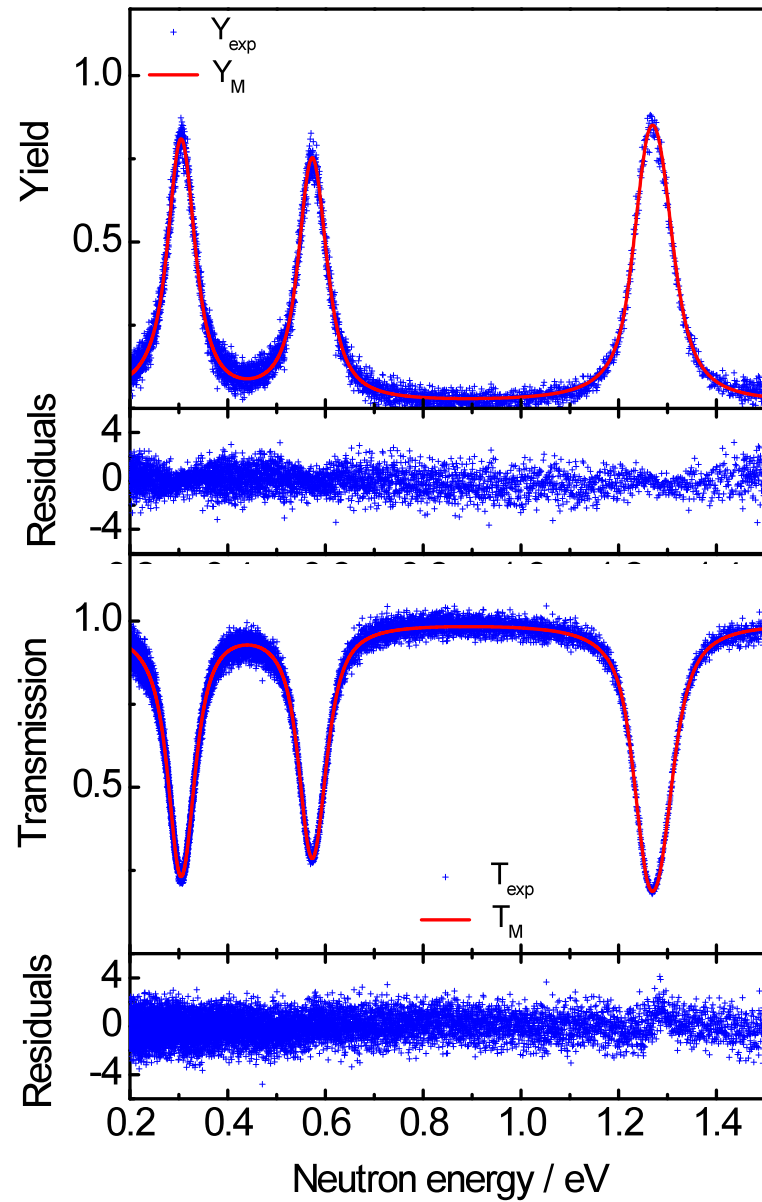
Sample properties: $^{242}\text{PuO}_2$ + carbon powder



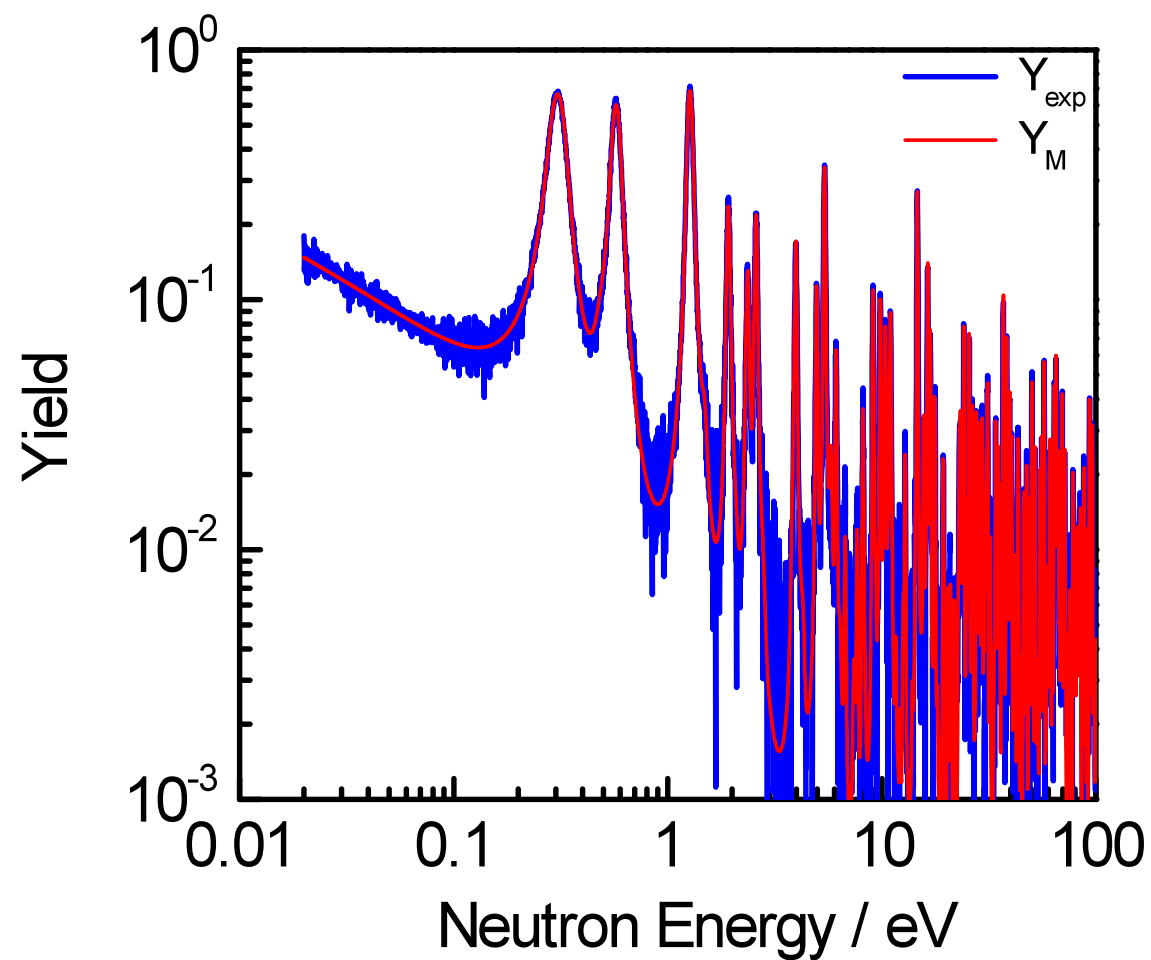
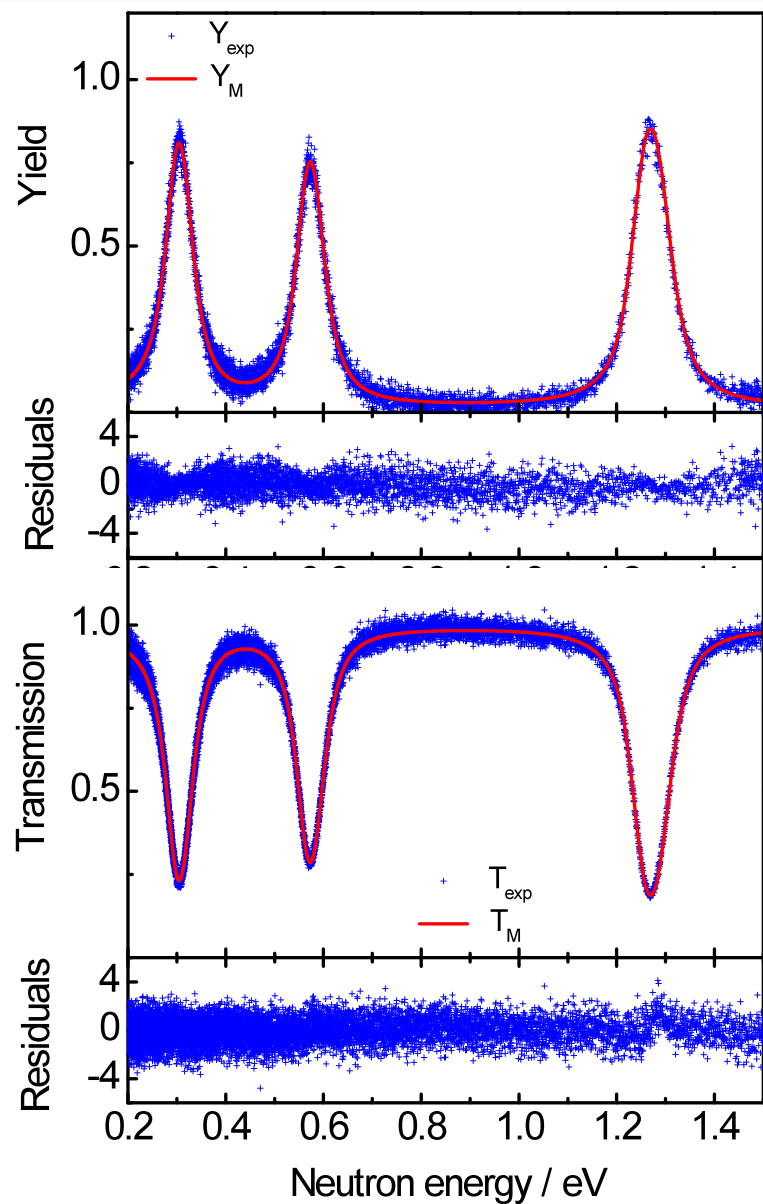
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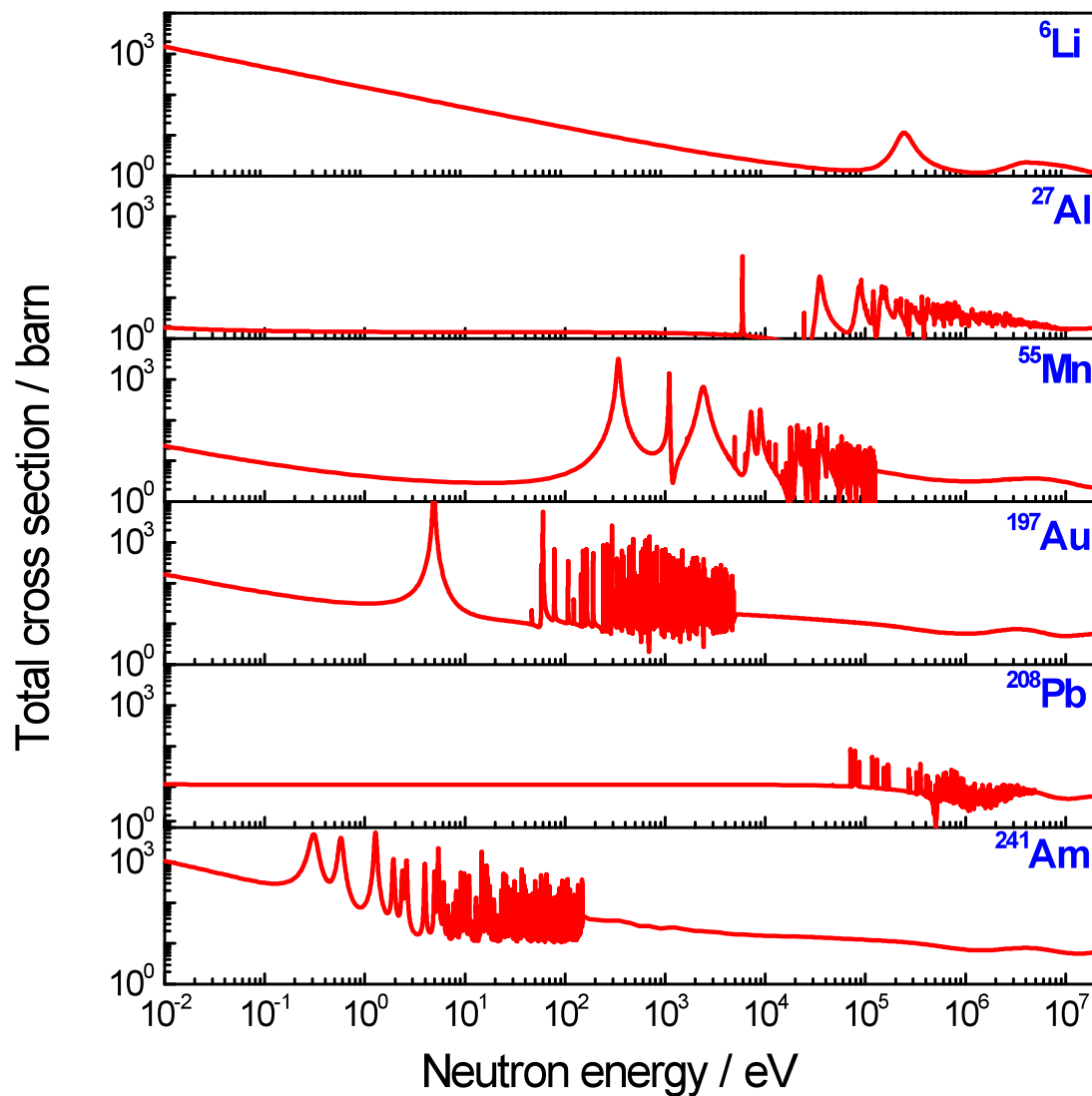
Resonance parameters for $^{241}\text{Am} + n$



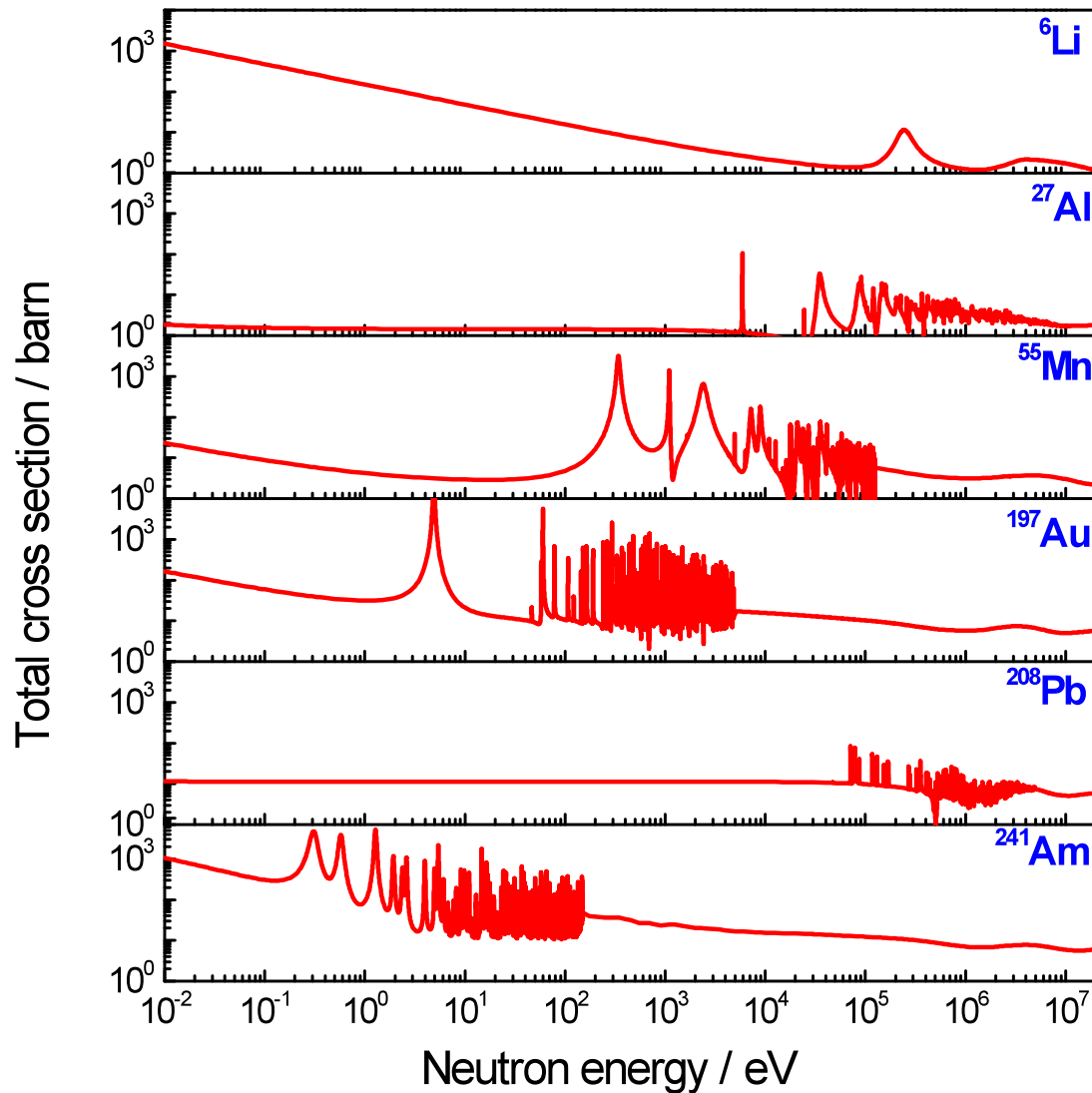
Resonance parameters for $^{241}\text{Am} + n$



Resonance structured cross sections

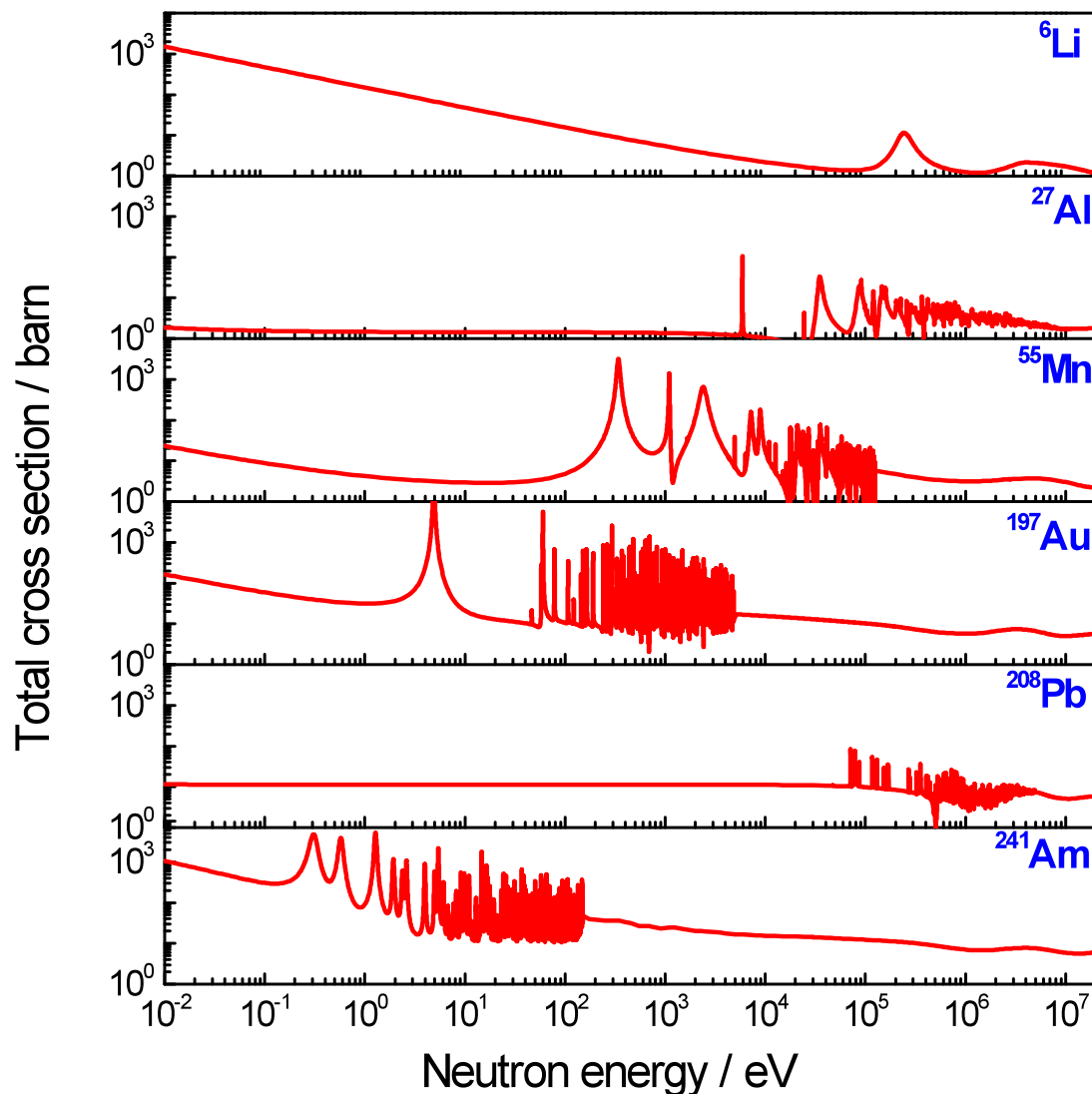


Resonance structured cross sections



Resonances appear at energies, which are specific for each nuclide

Neutron resonance analysis

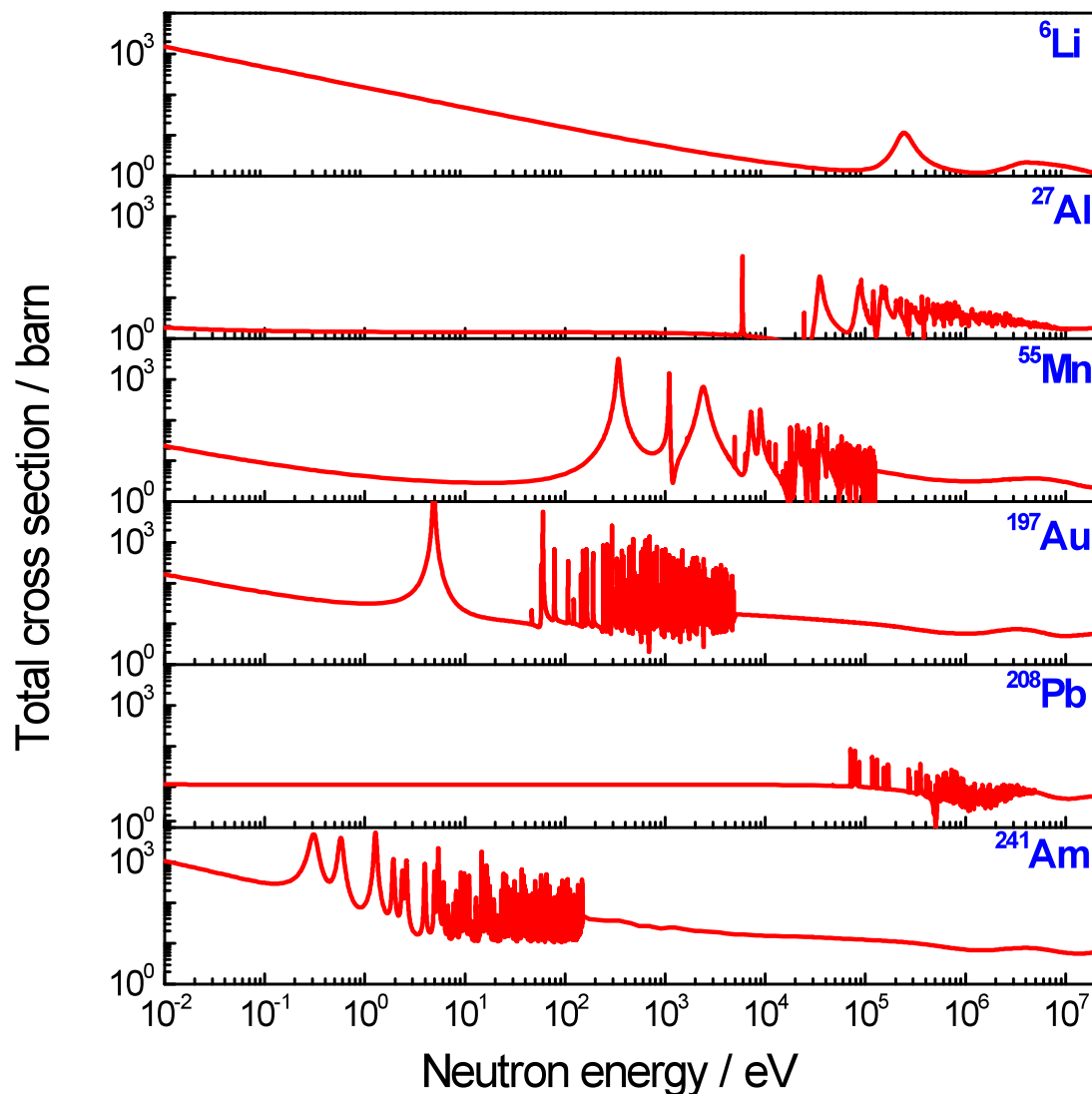


Resonances appear at energies, which are specific for each nuclide

▪ Resonances can be used to

- Identify and quantify nuclides
- Elemental (& isotopic) composition

NRCA & NRTA



Resonances appear at energies, which are specific for each nuclide

▪ Resonances can be used to

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▪ NRCA & NRTA

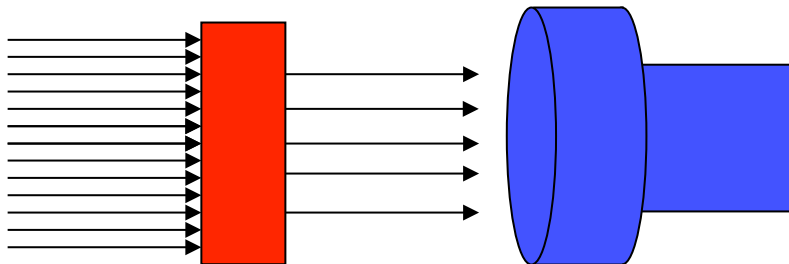
- Non - Destructive
- No sample preparation required
- Negligible residual activation

Cross section measurements

Transmission

$$T_{\text{exp}} = \frac{C_{\text{in}}}{C_{\text{out}}}$$

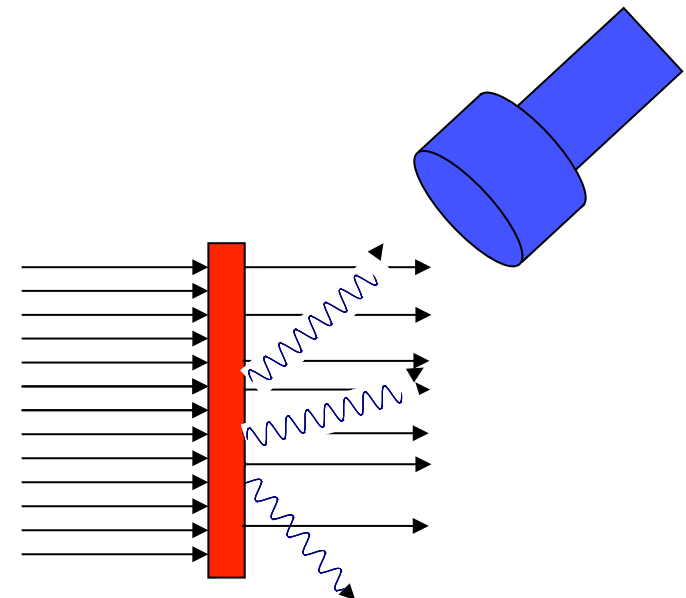
$$T \cong e^{-n \sigma_{\text{tot}}}$$



Capture

$$Y_{\gamma, \text{exp}} = \frac{C_{\gamma}}{\varepsilon_{\gamma} \varphi}$$

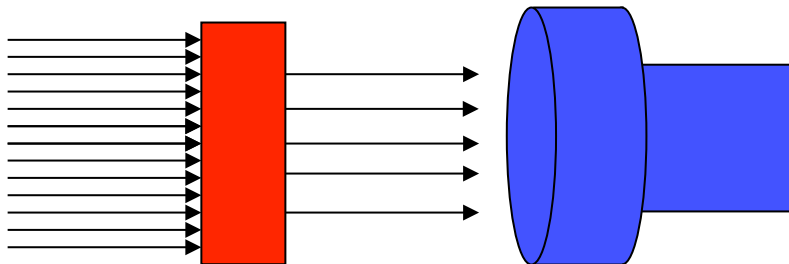
$$Y_{\gamma} \approx (1 - e^{-n \sigma_{\text{tot}}}) \frac{\sigma_{\gamma}}{\sigma_{\text{tot}}}$$



NRTA and NRCA

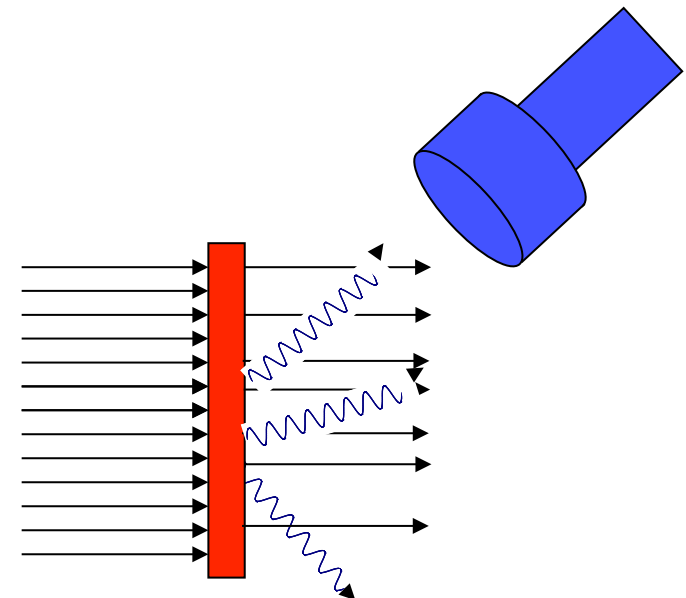
NRTA

$$T_{\text{exp}} = \frac{C_{\text{in}}}{C_{\text{out}}}$$
$$T \cong e^{-n \sigma_{\text{tot}}}$$



NRCA

$$Y_{\gamma, \text{exp}} = \frac{C_{\gamma}}{\varepsilon_{\gamma} \varphi}$$
$$Y_{\gamma} \approx (1 - e^{-n \sigma_{\text{tot}}}) \frac{\sigma_{\gamma}}{\sigma_{\text{tot}}}$$

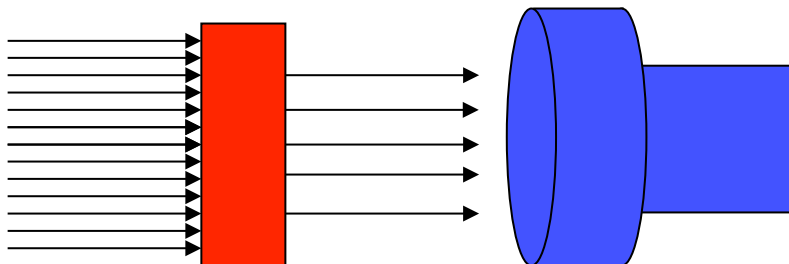


NRTA and NRCA

NRTA

$$T_{\text{exp}} = \frac{C_{\text{in}}}{C_{\text{out}}}$$
$$T \cong e^{-n \sigma_{\text{tot}}}$$

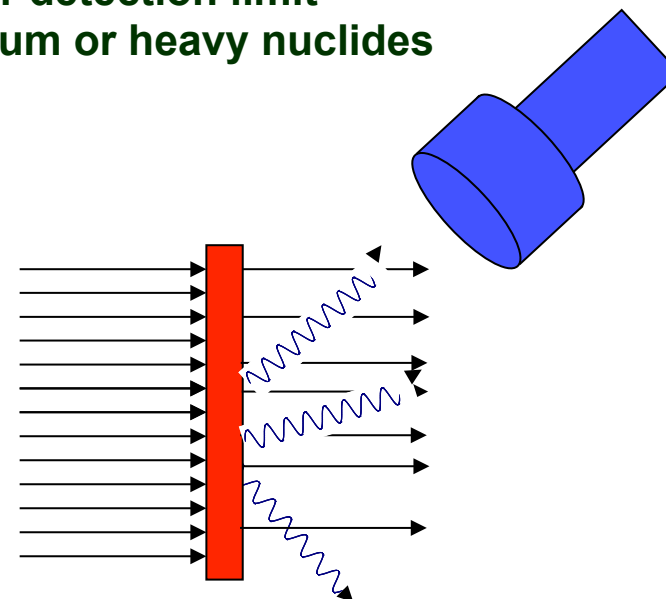
- + no calibration
- + radioactive samples
- + light nuclide or close to magic shells



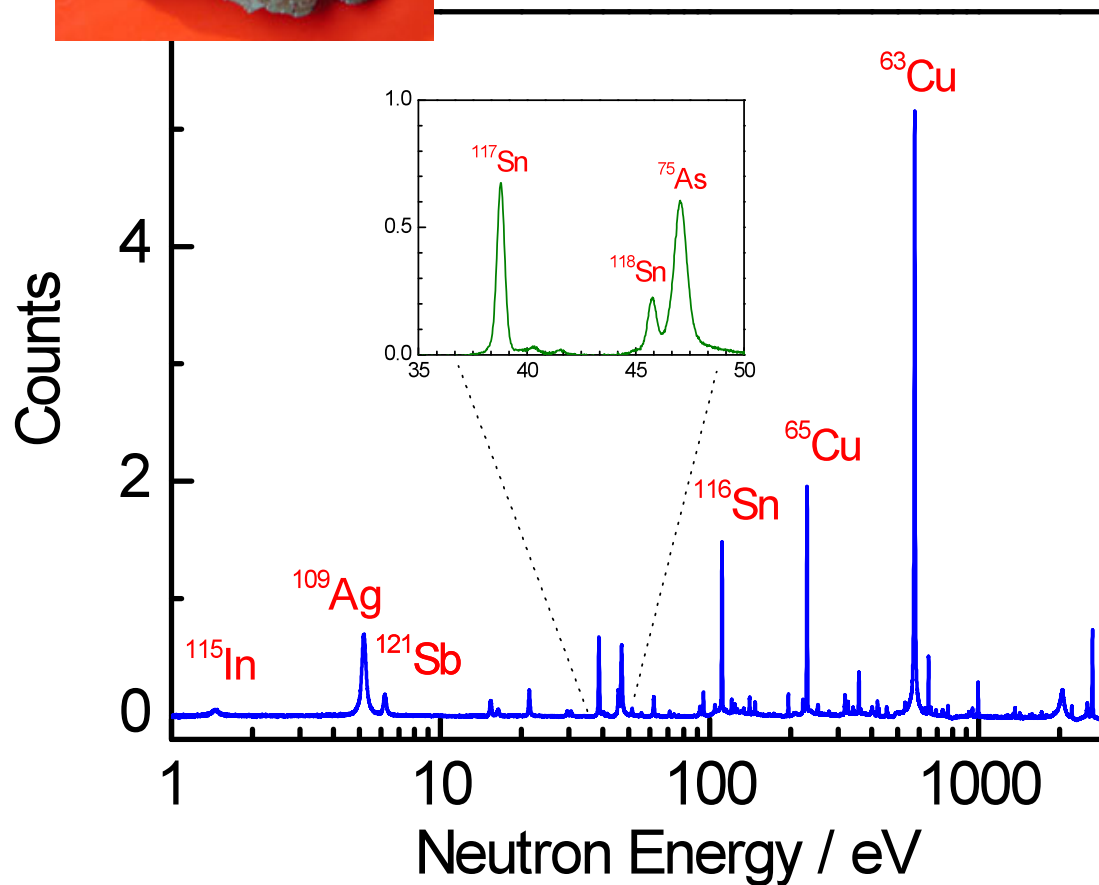
NRCA

$$Y_{\gamma, \text{exp}} = \frac{C_{\gamma}}{\varepsilon_{\gamma} \varphi}$$
$$Y_{\gamma} \approx (1 - e^{-n \sigma_{\text{tot}}}) \frac{\sigma_{\gamma}}{\sigma_{\text{tot}}}$$

- + better detection limit
- + medium or heavy nuclides



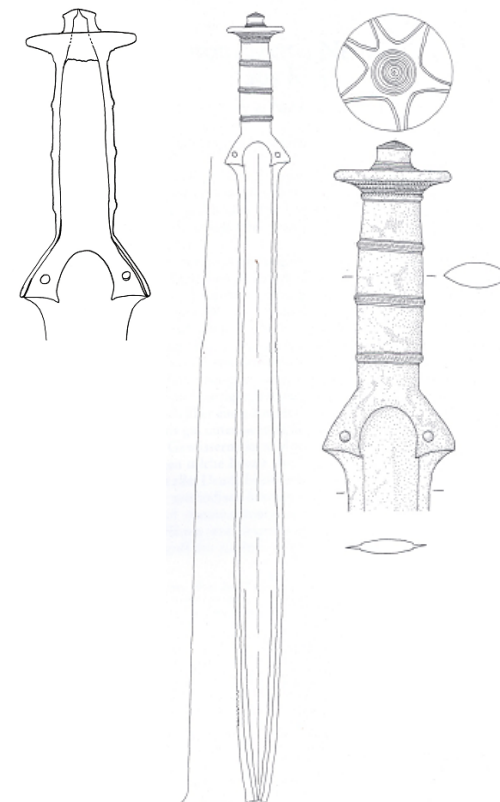
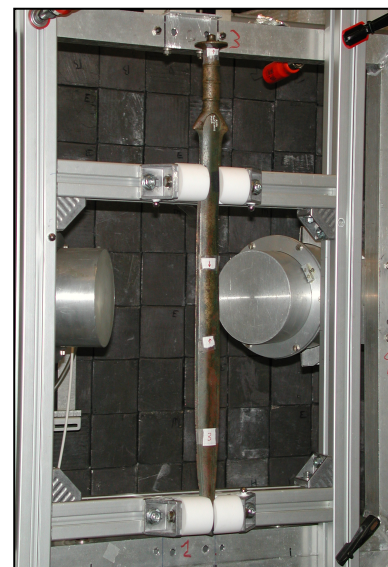
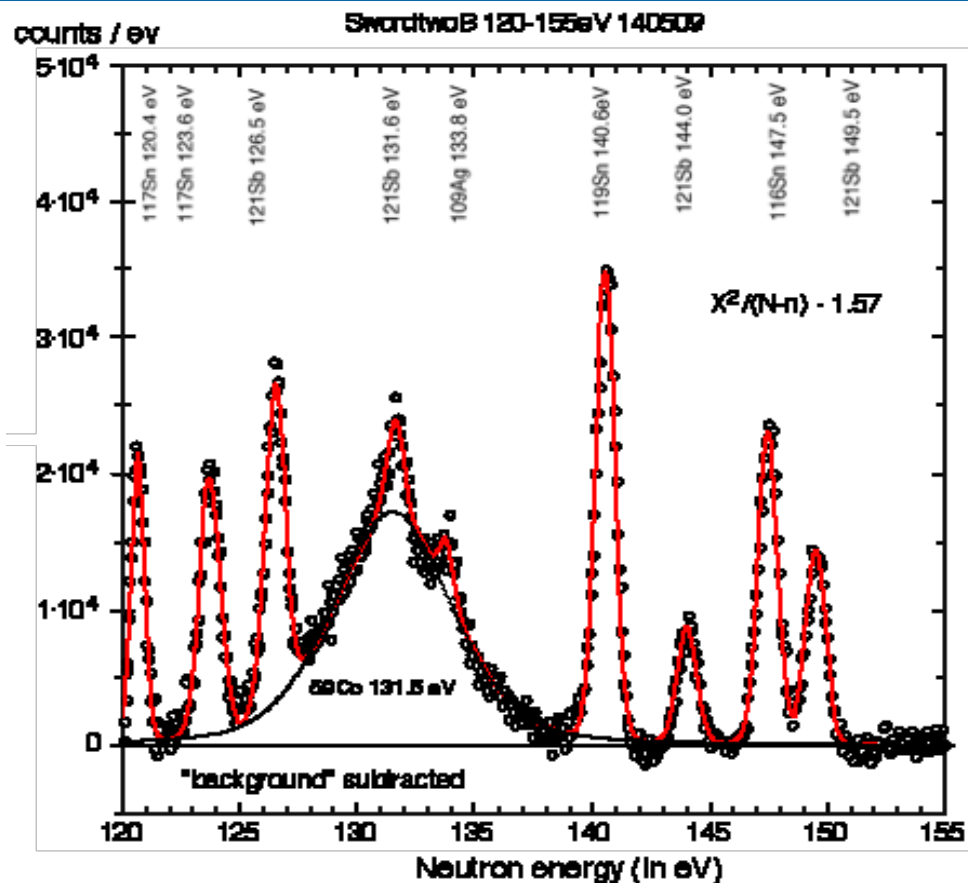
NRCA : example



Element	Fractions (%)	Isotope	Resonance (eV)
Cu	77.76 (0.11)	^{63}Cu	579.0
		^{65}Cu	230.0
Sn	20.85 (0.10)	^{112}Sn	94.8
		^{116}Sn	111.2
		^{117}Sn	38.8
		^{118}Sn	45.7
		^{119}Sn	222.6
		^{120}Sn	427.5
		^{122}Sn	1756.0
		^{124}Sn	62.0
As	0.34 (0.01)	^{75}As	47.0
Sb	0.196 (0.021)	^{121}Sb	6.24
		^{123}Sb	21.4
Ag	0.090 (0.01)	^{107}Ag	16.3
		^{109}Ag	5.2
Fe	0.770 (0.09)	^{56}Fe	1147.4
In	0.0061 (0.0003)	^{115}In	1.46

$m_{\text{NRCA}} = 13.0 (0.5) \text{ g}$
 $m_{\text{weight}} = 13.25 \text{ g}$

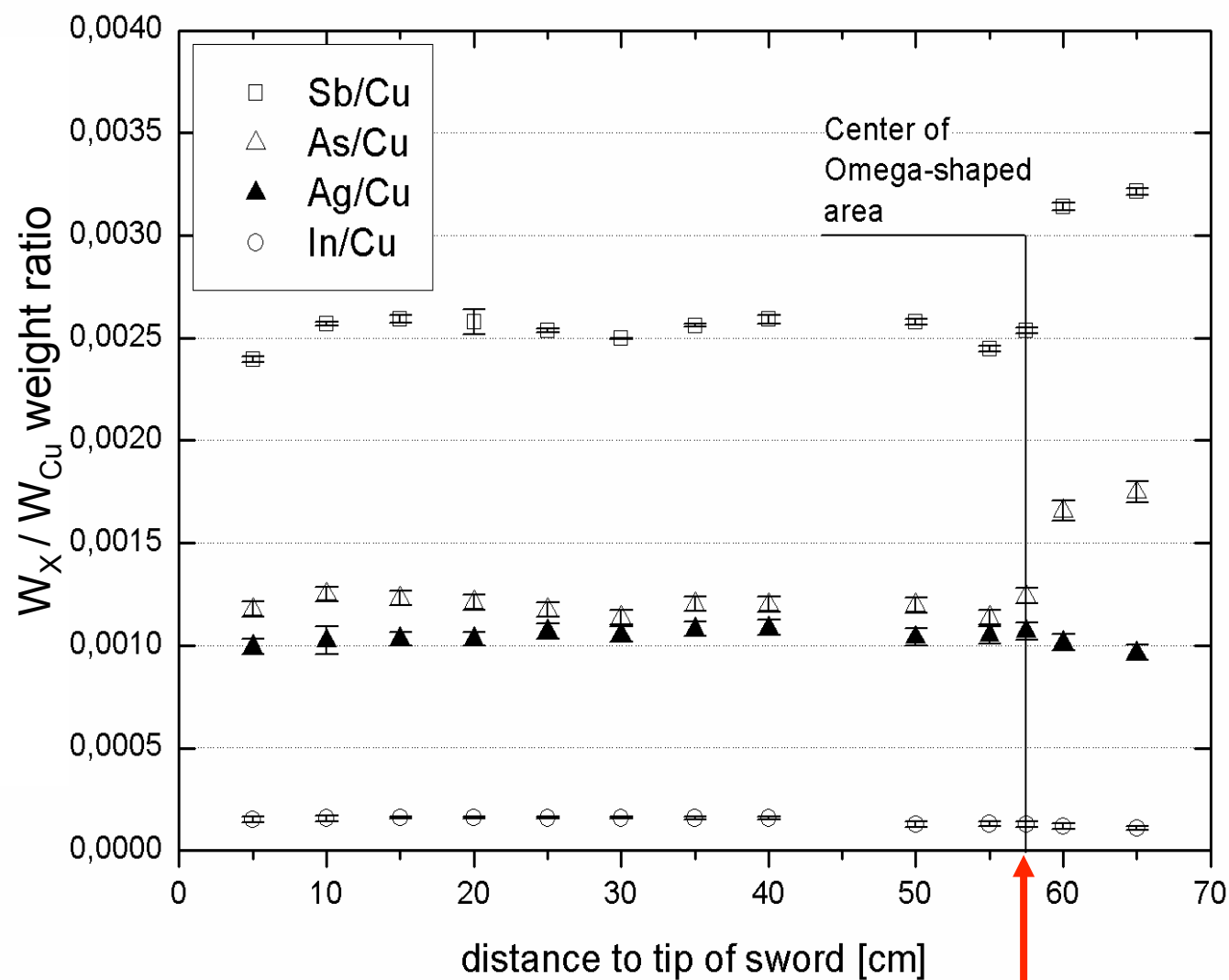
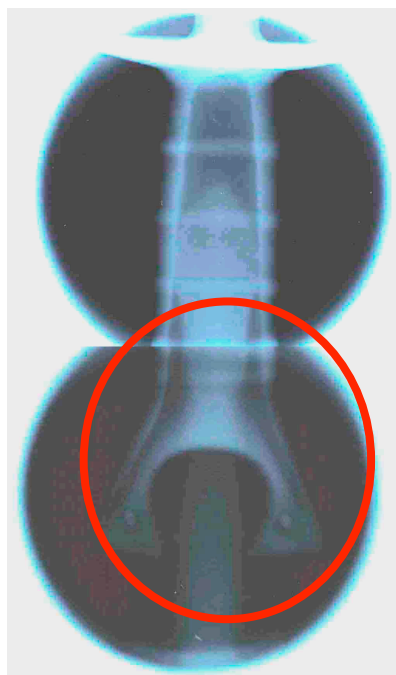
Bronze age sword



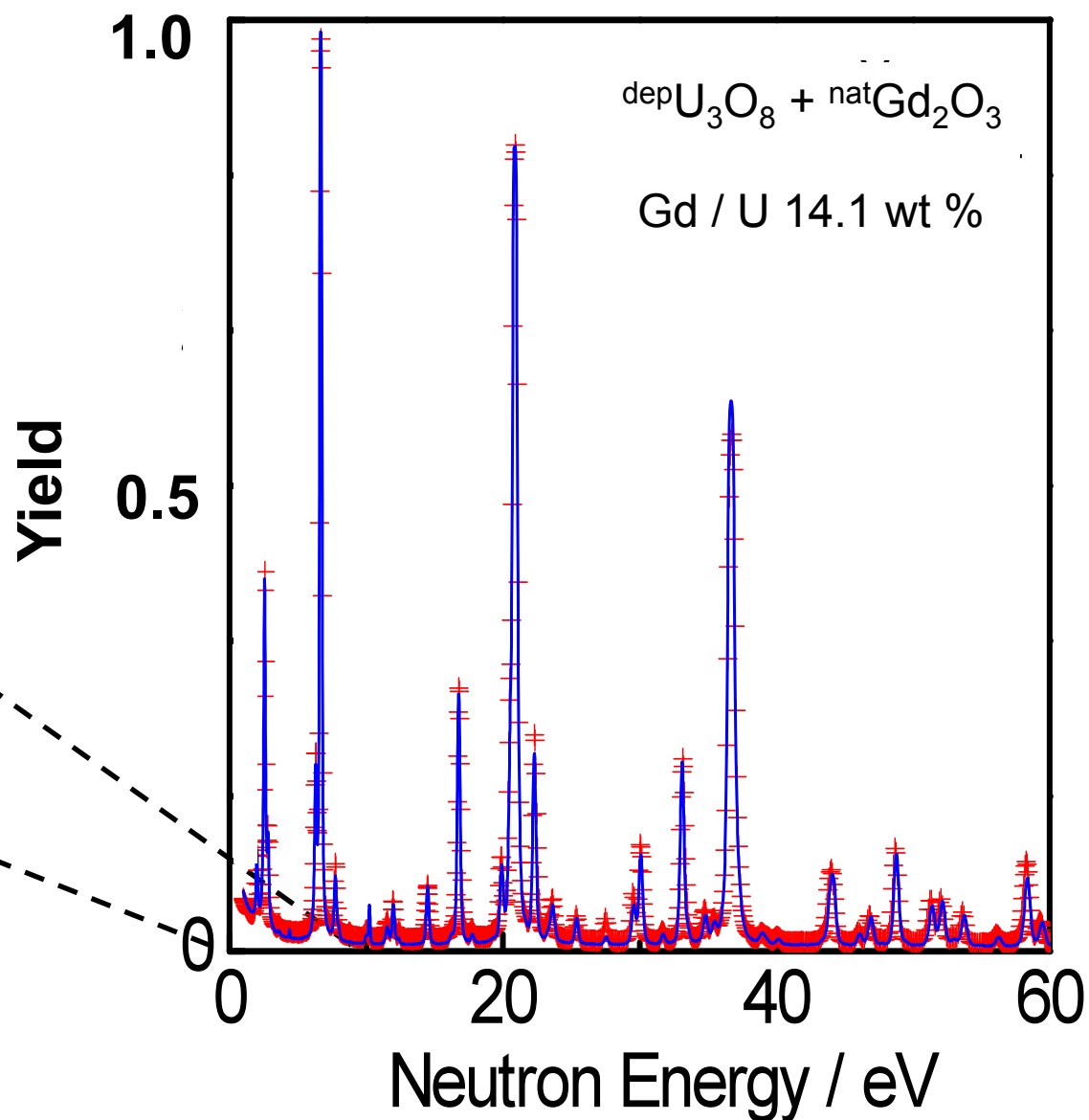
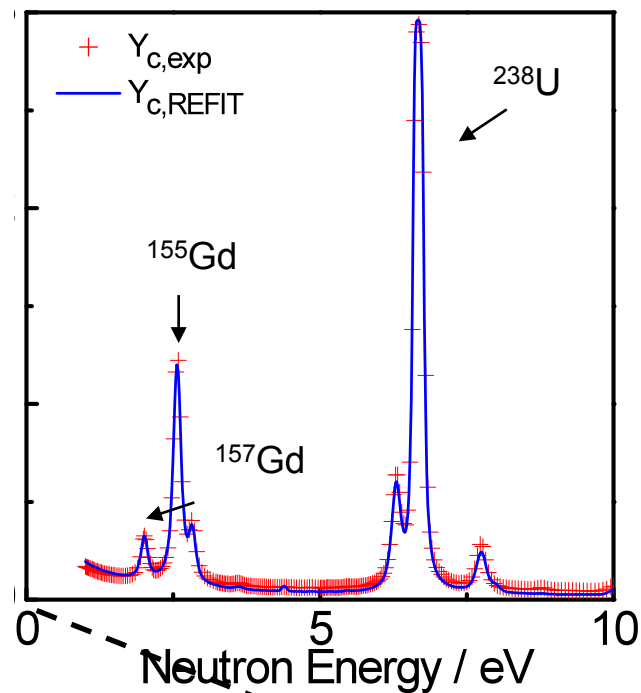
Tin – bronzes containing Sb, As and Ag

Presence of In and/or Co (not in all cases nor along all spots)

NRCA + γ -ray radiography

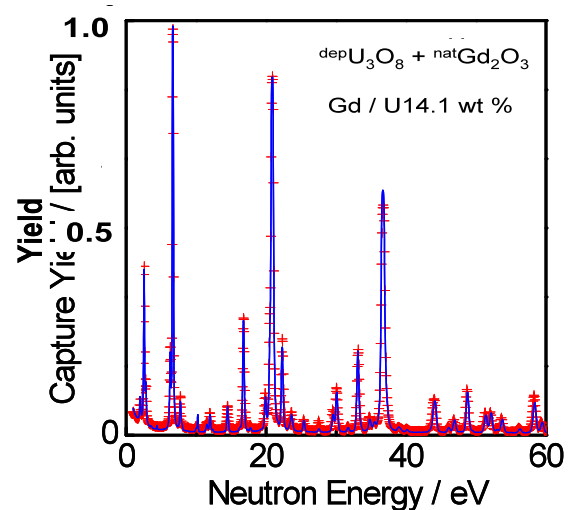


NRCA on nuclear materials



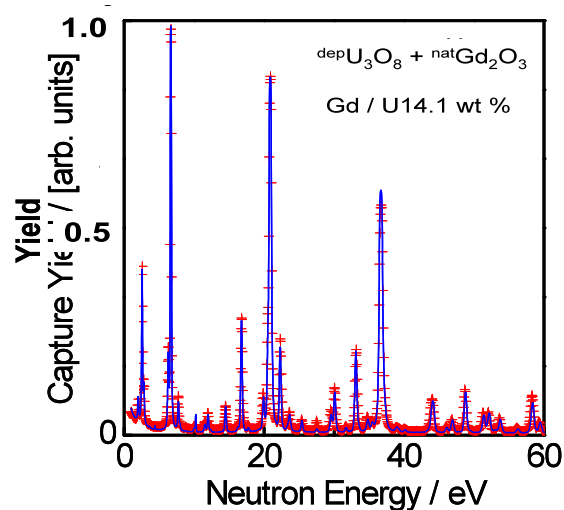
NRCA : RSA with REFIT

U / g	Gd / g	$n(^{151}\text{Gd}) / n(^{238}\text{U})$		$n(^{157}\text{Gd}) / n(^{238}\text{U})$	
		nat. abundance	NRCA	nat. abundance	NRCA
20.888	0.0636	$5.77 \cdot 10^{-4}$		$8.10 \cdot 10^{-4}$	
20.808	0.6208	$5.71 \cdot 10^{-3}$		$8.03 \cdot 10^{-4}$	
18.858	2.8240	$3.13 \cdot 10^{-2}$		$3.36 \cdot 10^{-3}$	



NRCA : RSA with REFIT

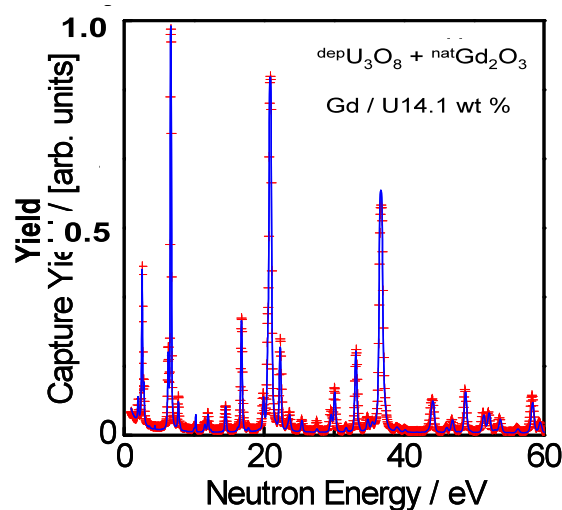
U / g	Gd / g	$n(^{151}\text{Gd}) / n(^{238}\text{U})$		$n(^{157}\text{Gd}) / n(^{238}\text{U})$	
		nat. abundance	NRCA	nat. abundance	NRCA
20.888	0.0636	$5.77 \cdot 10^{-4}$	$(5.78 \pm 0.04) \cdot 10^{-4}$	$8.10 \cdot 10^{-4}$	
20.808	0.6208	$5.71 \cdot 10^{-4}$	$(5.73 \pm 0.01) \cdot 10^{-4}$	$8.03 \cdot 10^{-4}$	
18.858	2.8240	$3.13 \cdot 10^{-2}$	$(3.14 \pm 0.01) \cdot 10^{-2}$	$3.36 \cdot 10^{-2}$	



NRCA : RSA with REFIT



U / g	Gd / g	$n(^{151}\text{Gd}) / n(^{238}\text{U})$		$n(^{157}\text{Gd}) / n(^{238}\text{U})$	
		nat. abundance	NRCA	nat. abundance	NRCA
20.888	0.0636	$5.77 \cdot 10^{-4}$	$(5.78 \pm 0.04) \cdot 10^{-4}$	$8.10 \cdot 10^{-4}$	$(8.59 \pm 0.07) \cdot 10^{-4}$
20.608	0.6208	$5.71 \cdot 10^{-3}$	$(5.73 \pm 0.01) \cdot 10^{-3}$	$8.03 \cdot 10^{-3}$	$(8.53 \pm 0.02) \cdot 10^{-3}$
18.858	2.8240	$3.13 \cdot 10^{-2}$	$(3.14 \pm 0.01) \cdot 10^{-2}$	$3.36 \cdot 10^{-2}$	$(3.51 \pm 0.03) \cdot 10^{-2}$

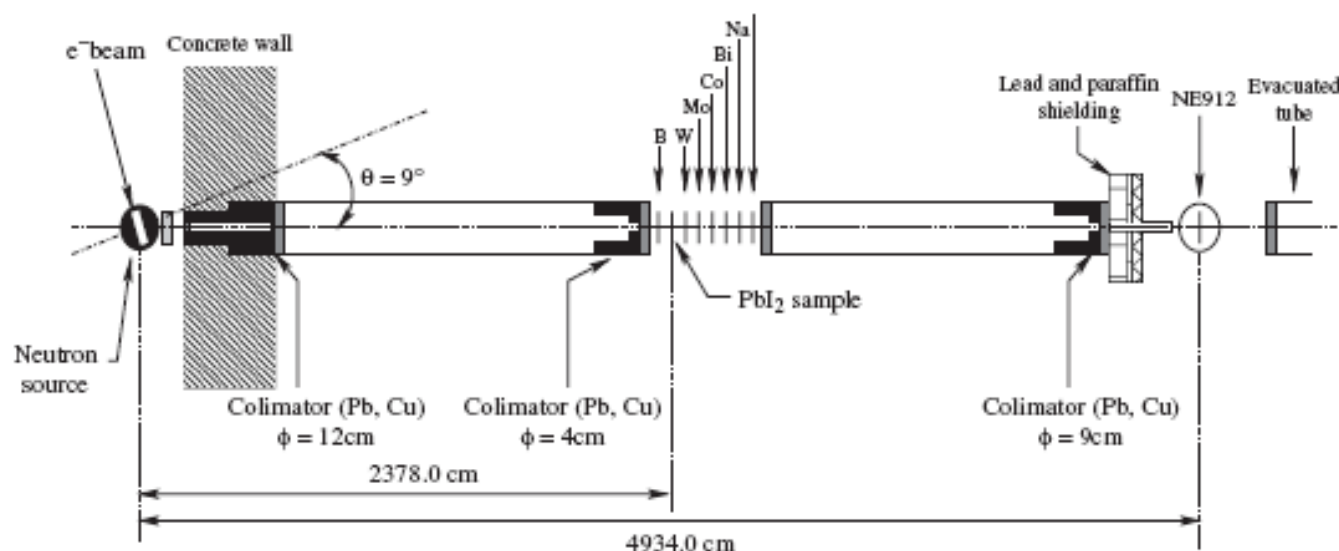


Characterisation of PbI_2 sample by NRTA

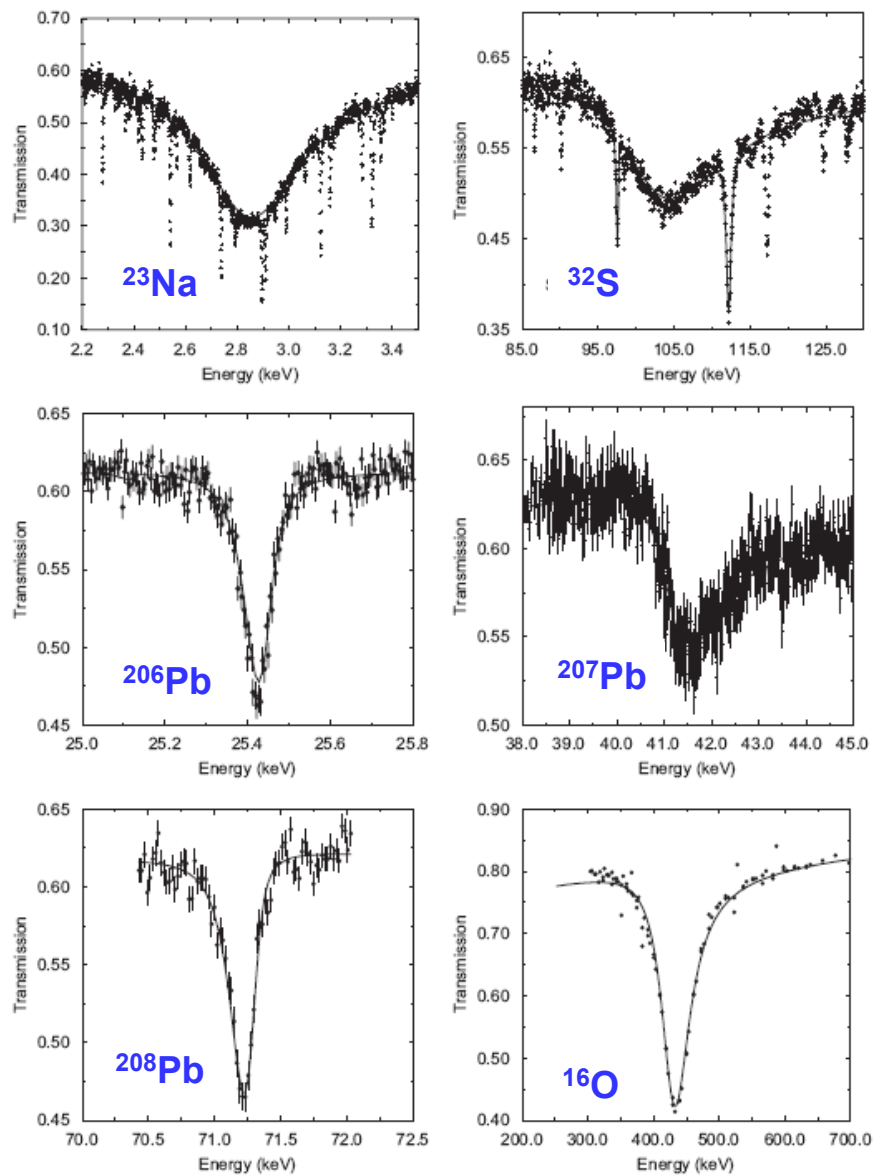
- **IRMM sample preparation group extracted** (NIMA 480 (2002) 204) :
 - 150 g Iodine (powder) from 210 liter reprocessed waste
 - (1.3 g/l Iodine and 40 MBq/l)
- **Sample characterisation: by mass spectrometry , (N)AA and NRTA**



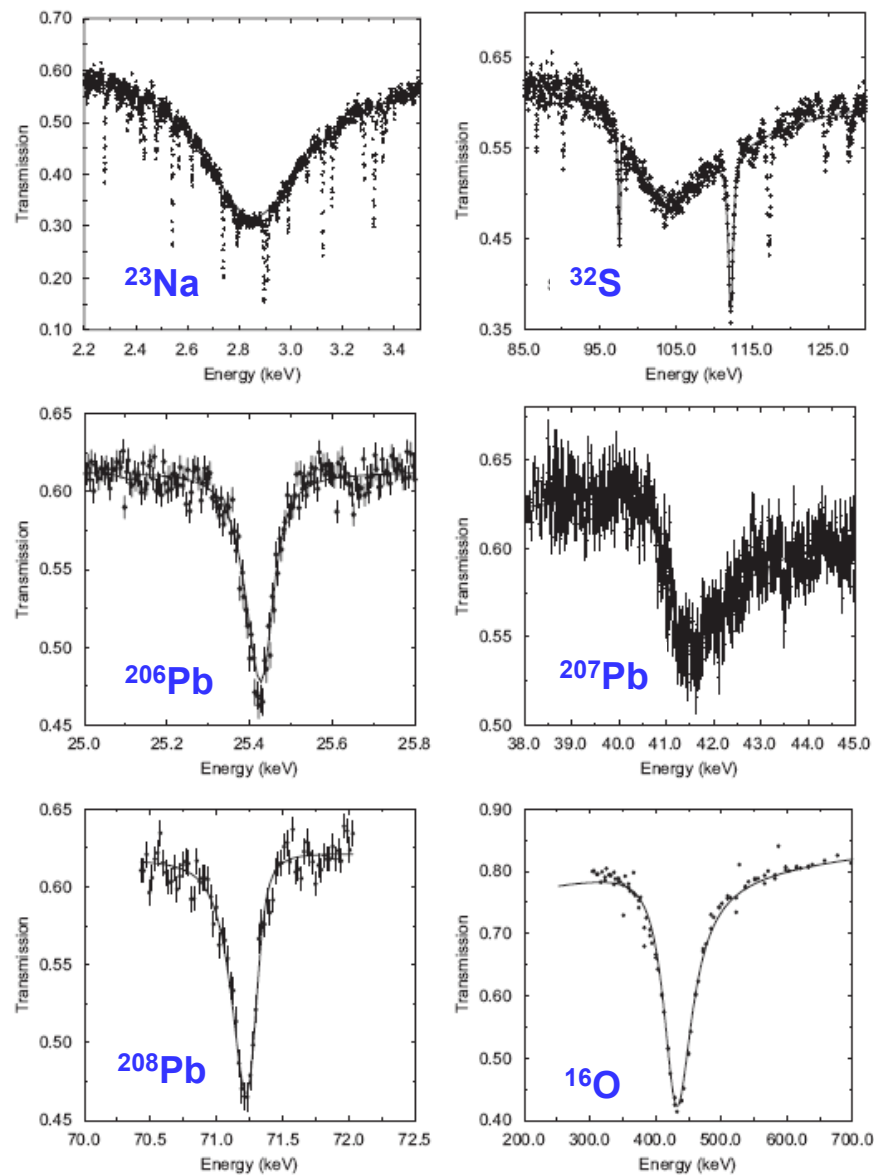
Fig. 3. Addition of $\text{Pb}(\text{NO}_3)_2$ to the iodide solution to precipitate lead iodide.



Characterisation of PbI_2 by NRTA

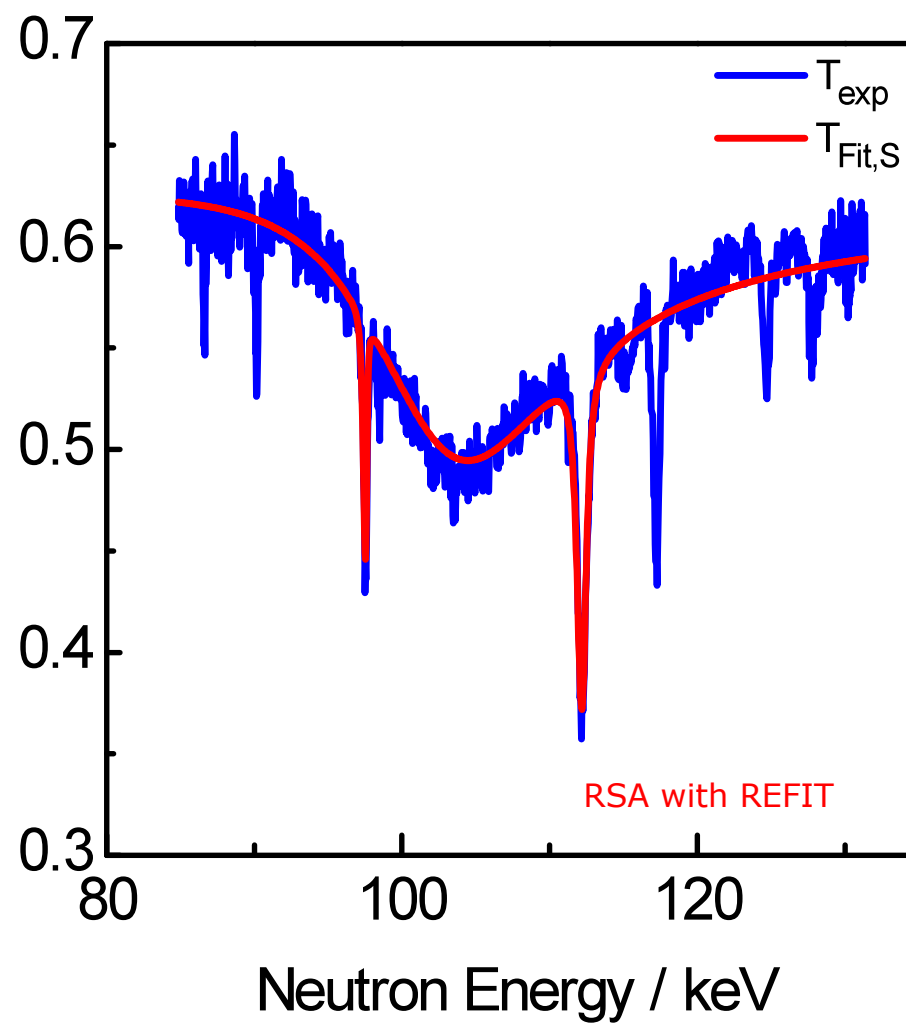


Characterisation of PbI_2 by NRTA



Transmission

RSA with REFIT



Advantages of NRTA



Element		NRTA (GELINA)		NAA		Mass spectrometry (PSI)	
Iodine	total	20.24	(0.41)	19.75	(0.61)	19.86	(0.41)
	¹²⁷ I	3.44	(0.05)	3.35	(0.10)	3.36	(0.08)
	¹²⁹ I	16.80	(0.40)	16.40	(0.60)	16.50	(0.40)
Lead	total	52.30	(1.70)	51.10	(1.80)		
	²⁰⁶ Pb	12.80	(0.50)				
	²⁰⁷ Pb	11.50	(0.10)				
	²⁰⁸ Pb	27.10	(1.70)				
Sulfur		5.44	(0.03)				
Sodium		0.72	(0.02)			1.00	(0.15)
Oxygen		13.92	(0.05)			14.50	(1.50)
Hydrogen		< 0.13				0.02	(0.002)
Nitrogen						1.20	(0.40)

Almost all elements present in the sample have been analyzed
Isotopic composition of Pb

From Analysis to Imaging

GELINA at IRMM



ISIS at RAL



PSND for imaging at ISIS

10 x 10 pixelated neutron detector, 100 ^6Li -glass scintillators ($2 \times 2 \times 8 \text{ mm}^3$)

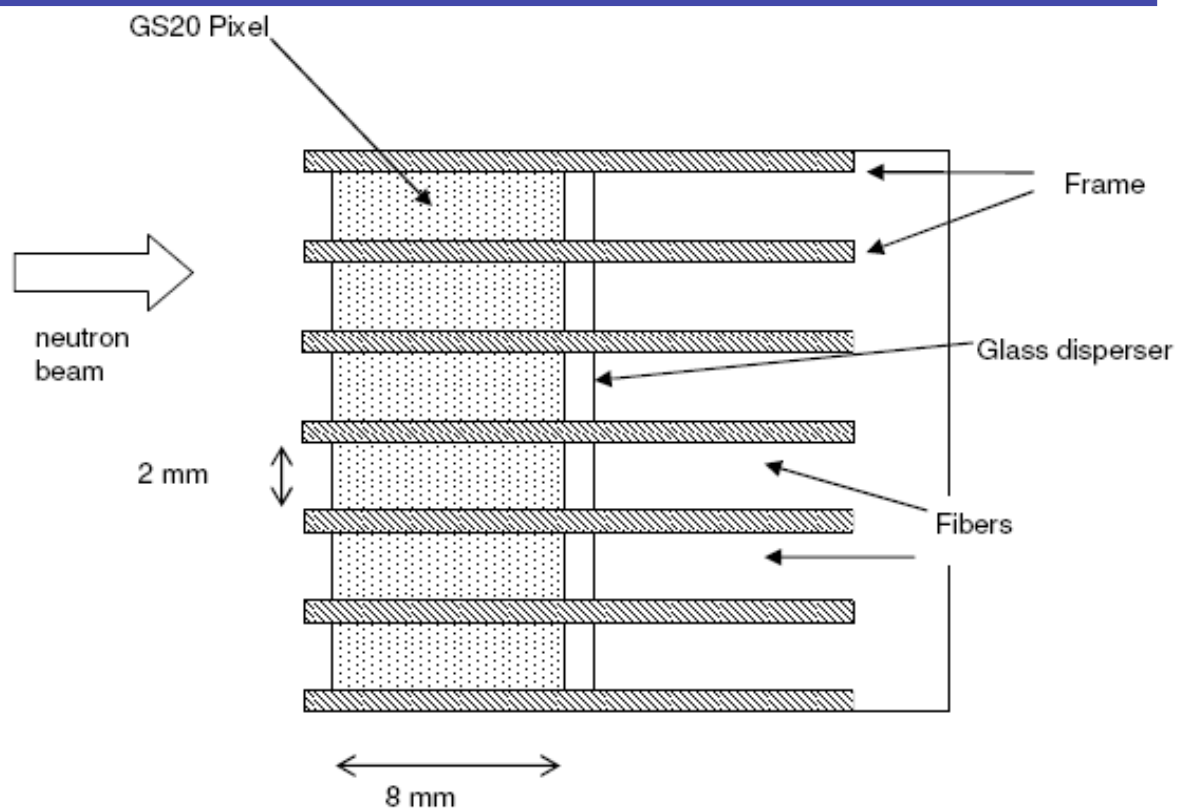
7 position sensitive 16 channel PMT



Detection efficiency

25 % at 10 eV

4 % at 1 keV



Test of PSND for NRTI at GELINA



Tested for

- Time resolution
- Cross talk
- Procedure to determine T_{exp} and deduce n in non-ideal geometry

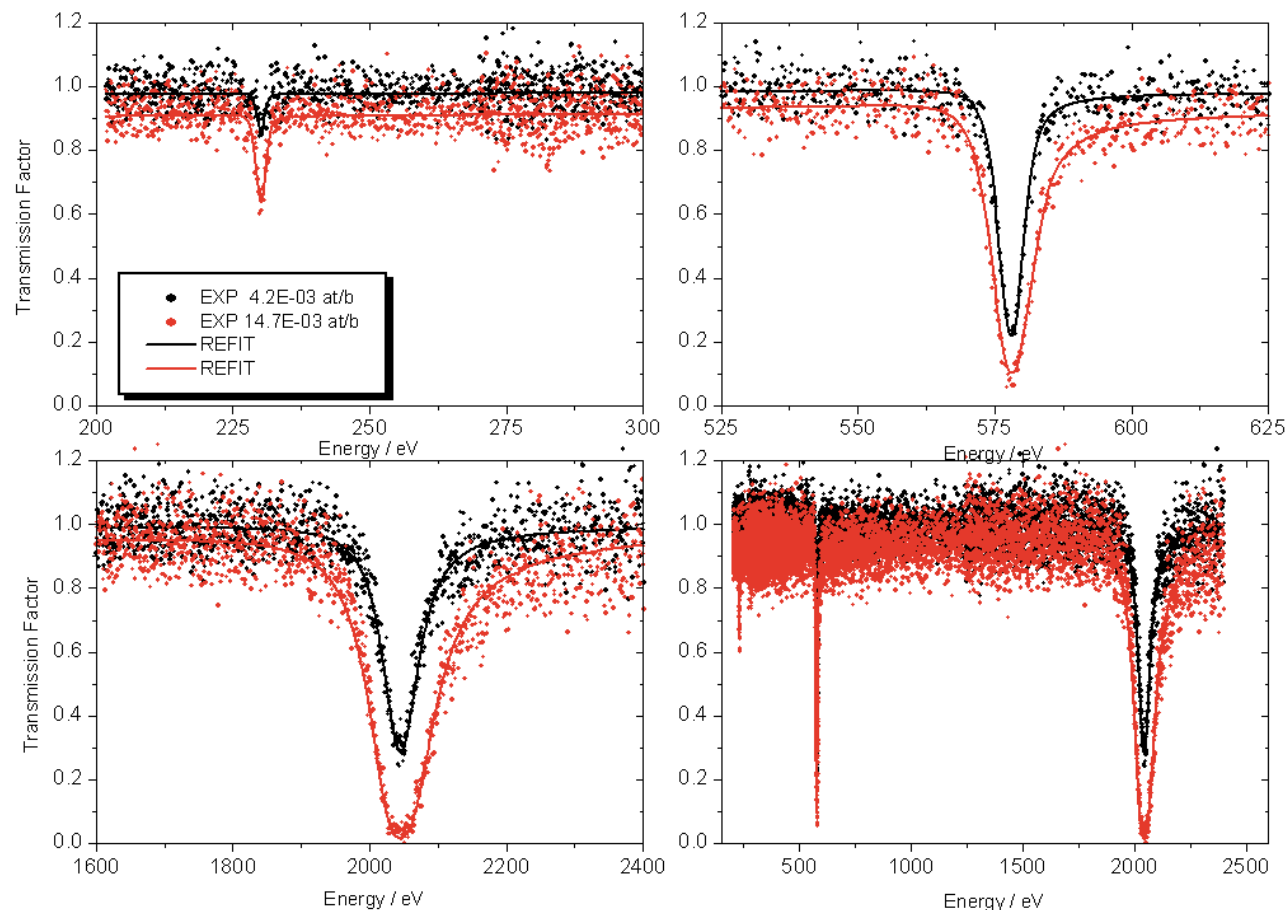


Test of PSND for NRTI at GELINA



Tested for

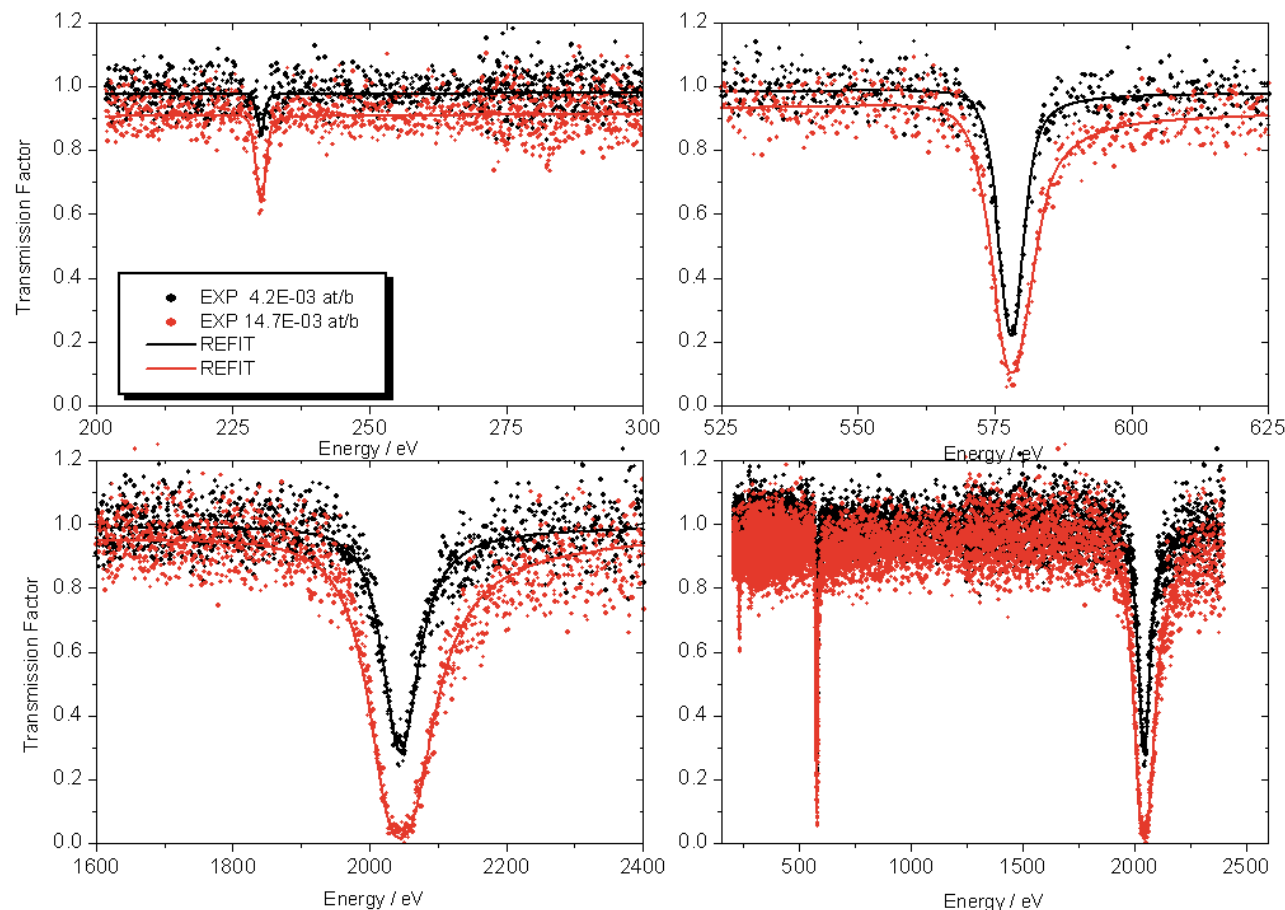
- Time resolution
 - Cross talk
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- Measurements on Cu-discs



Test of PSND for NRTI at GELINA

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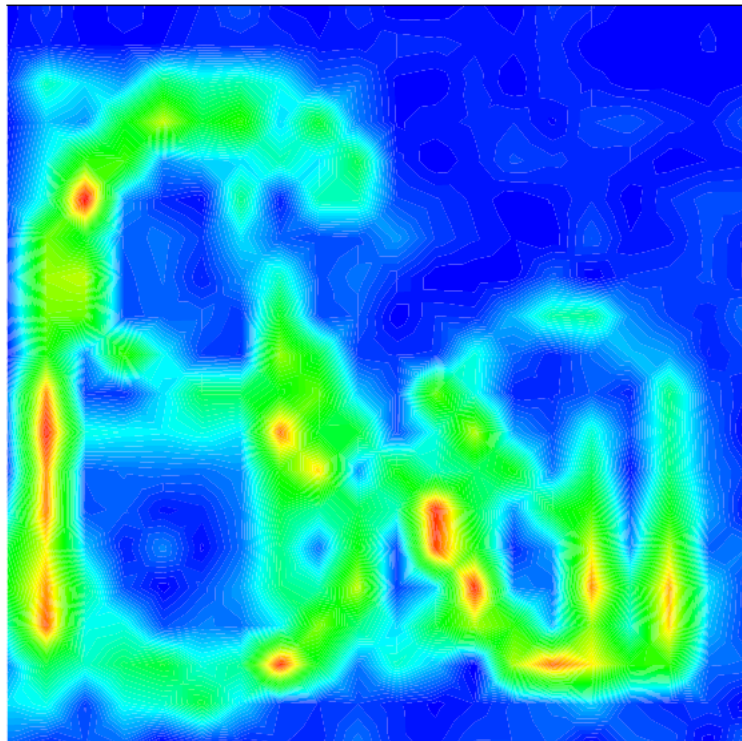
NRTA with REFIT

$$n_{\text{Cu}} = (4.29 \pm 0.05) \cdot 10^{-3} \text{ at/b} \Leftrightarrow 4.22 \cdot 10^{-3} \text{ at/b}$$

$$n_{\text{Cu}} = (1.49 \pm 0.01) \cdot 10^{-2} \text{ at/b} \Leftrightarrow 1.47 \cdot 10^{-2} \text{ at/b}$$

NRTI with PSND at ISIS

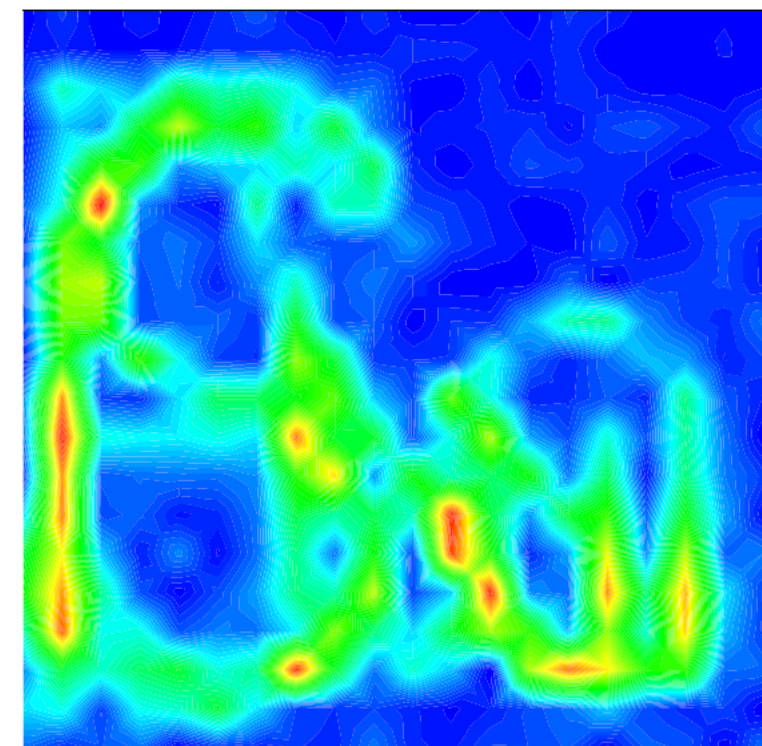
Total counts in TOF spectrum



40 mm

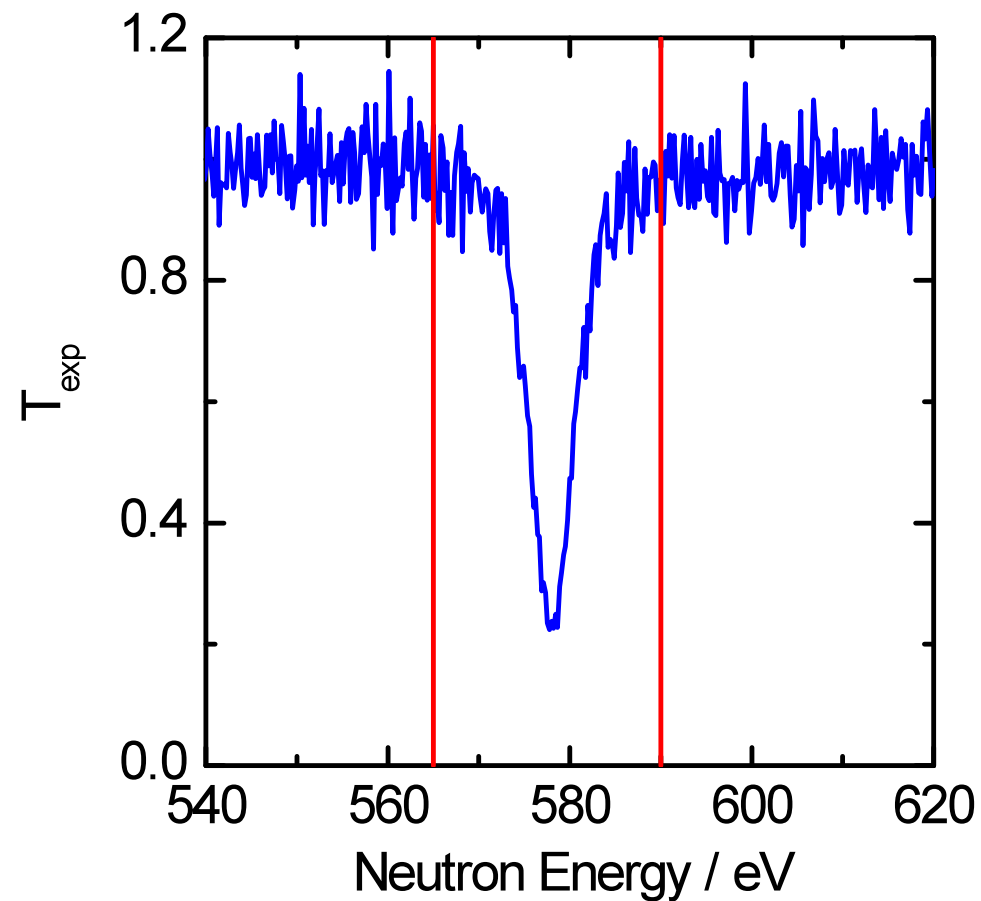
NRTI with PSND at ISIS

Total counts in TOF spectrum



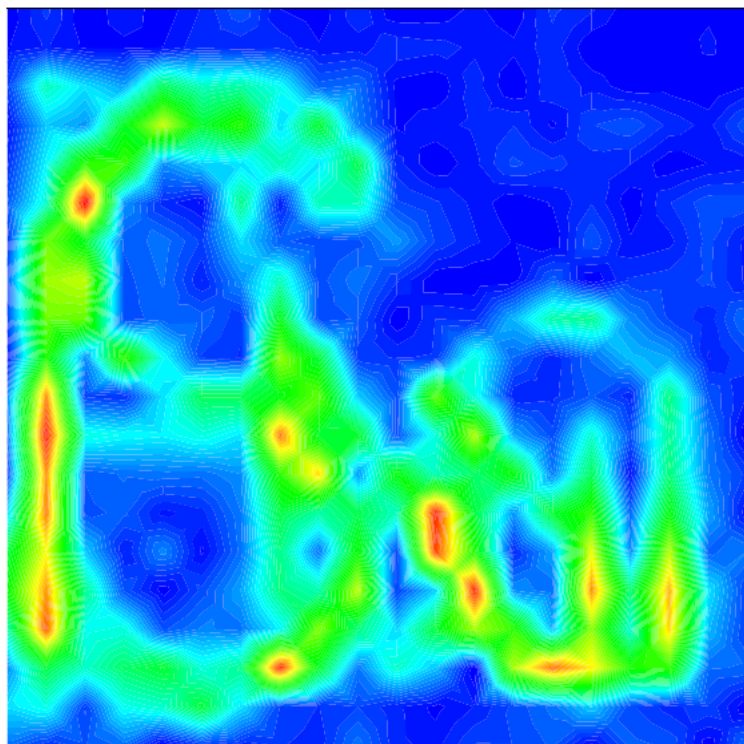
40 mm

Select region : $E_{\mu} = 230$ eV for $^{65}\text{Cu} + n$



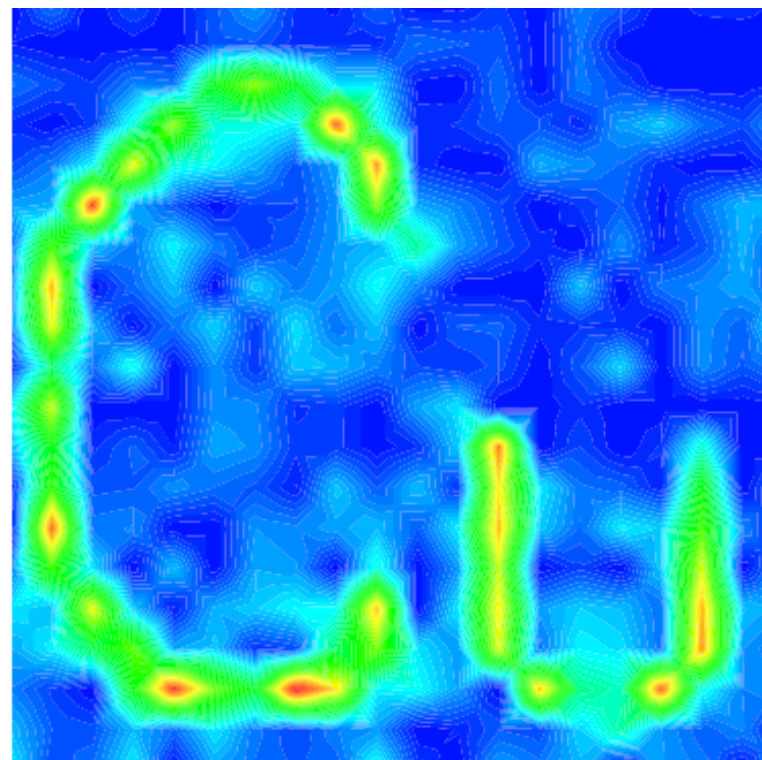
NRTI with PSND at ISIS

Total counts in TOF spectrum



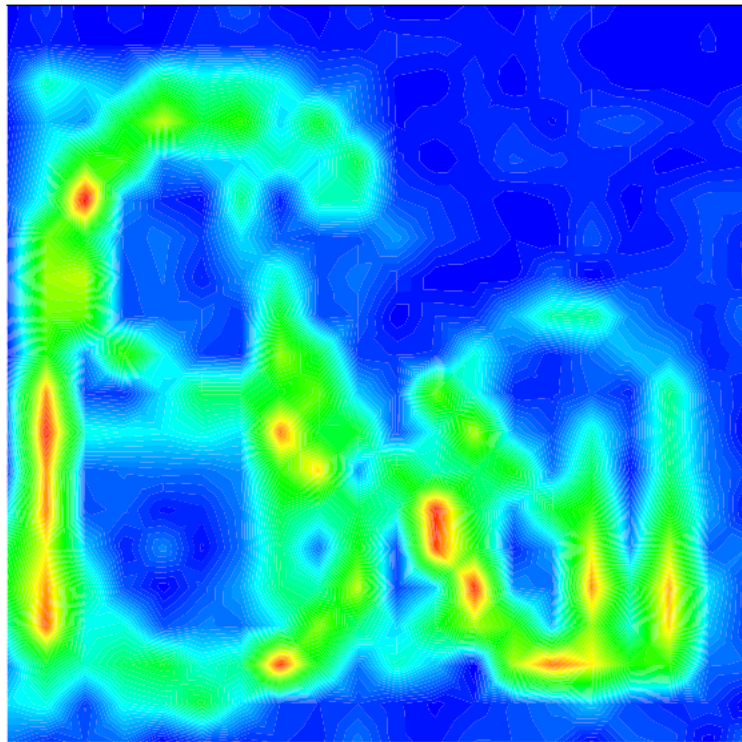
40 mm

Select region : $E_{\mu} = 230$ eV for $^{65}\text{Cu} + n$



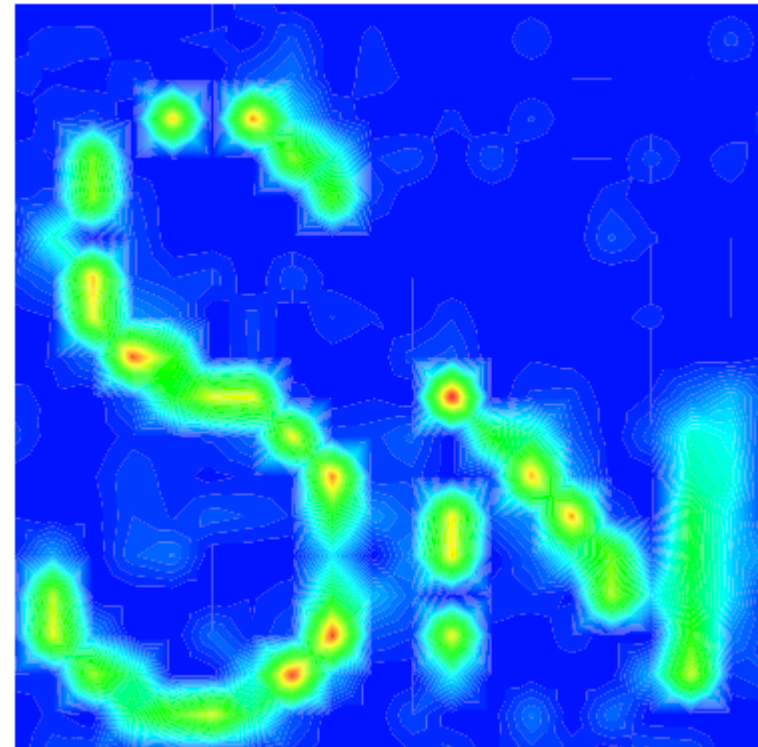
NRTI with PSND at ISIS

Total counts in TOF spectrum



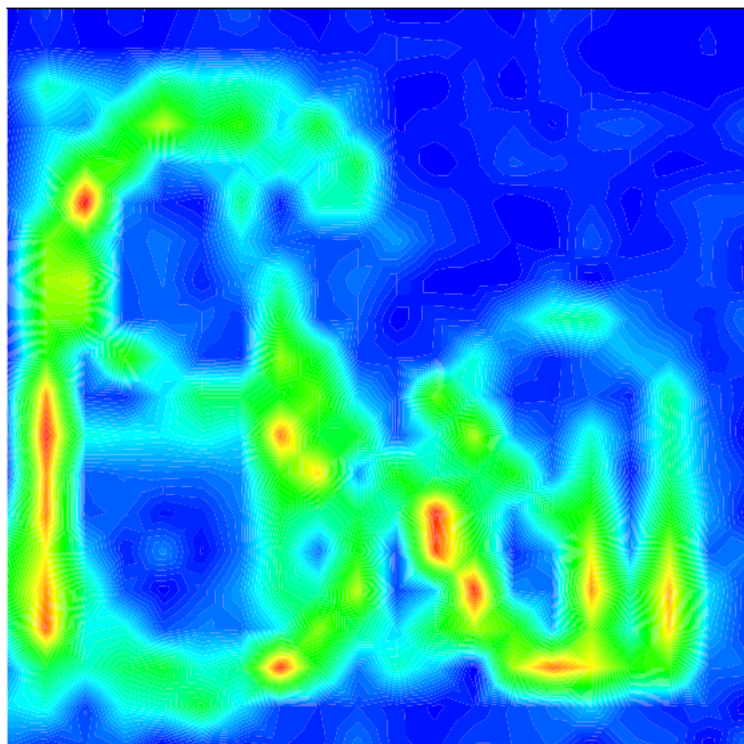
40 mm

Select region : $E_{\mu} = 111.2$ eV for $^{116}\text{Sn} + n$



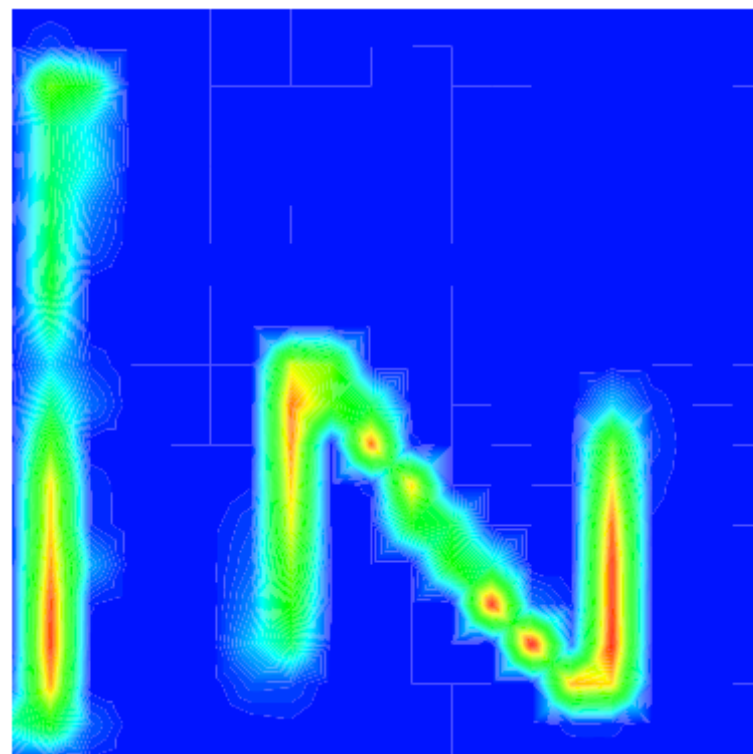
NRTI with PSND at ISIS

Total counts in TOF spectrum

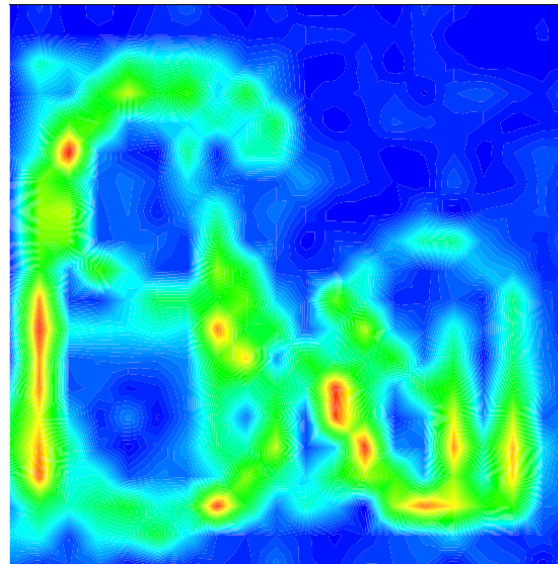


40 mm

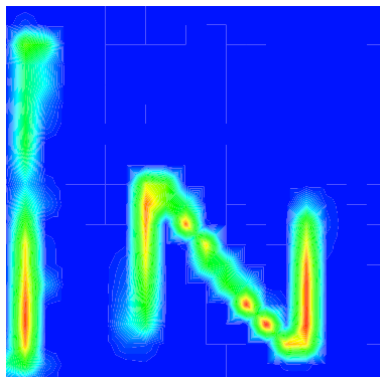
Select region : $E_{\mu} = 1.46$ eV for $^{115}\text{In} + n$



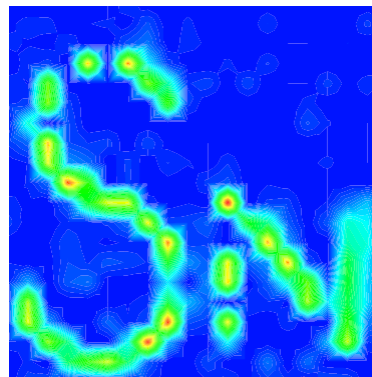
NRTI with PSND at ISIS



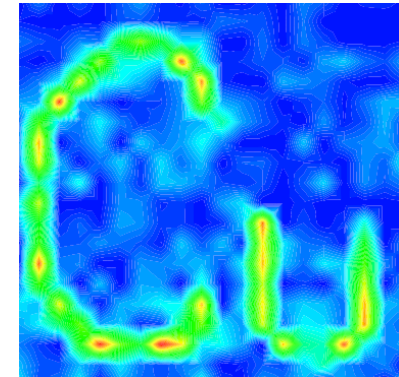
40 mm



$E_n = 1.46 \text{ eV}$



$E_n = 111.2 \text{ eV}$



$E_n = 230 \text{ eV}$

NRTI with PSND at ISIS

10 x 10 pixelated neutron detector, 100 ^6Li -glass scintillators (2 x 2 x 9 mm³)

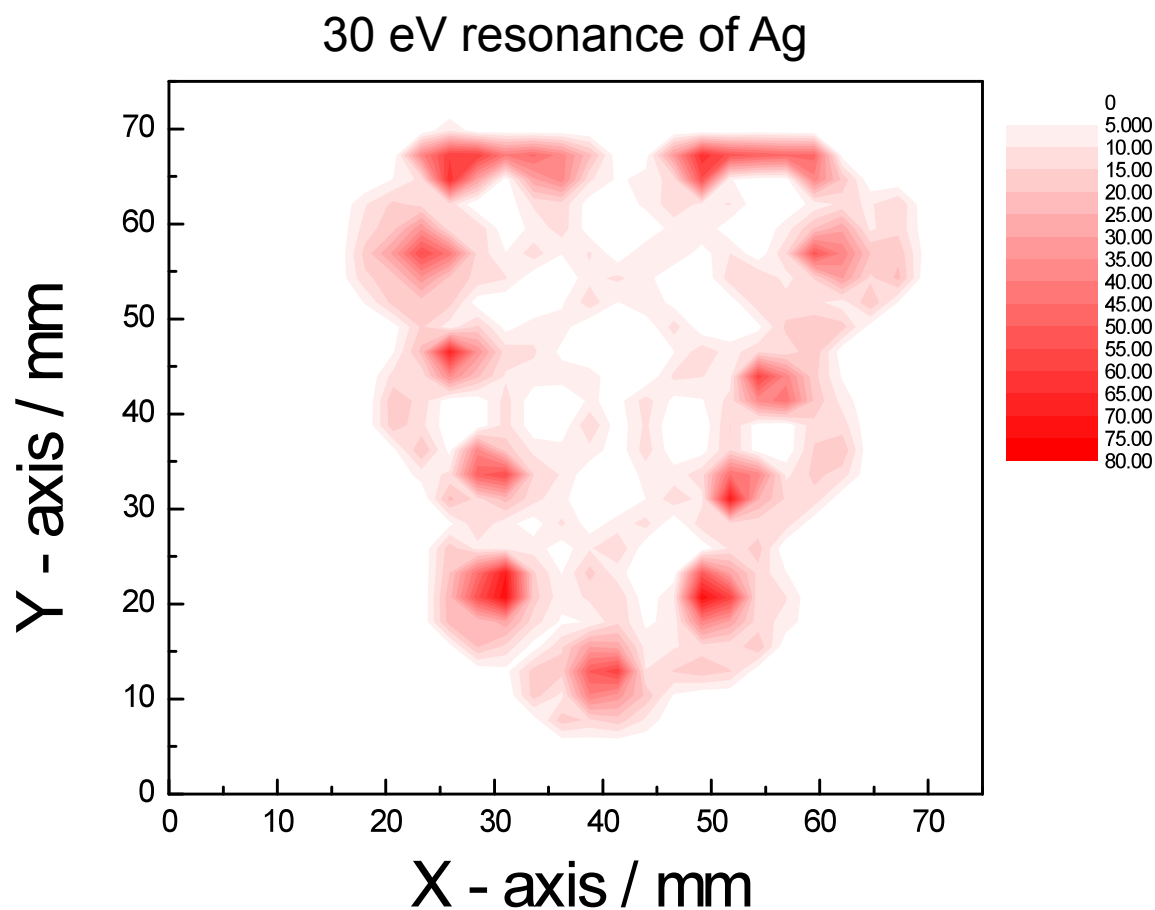
7 position sensitive 16 channel PMT



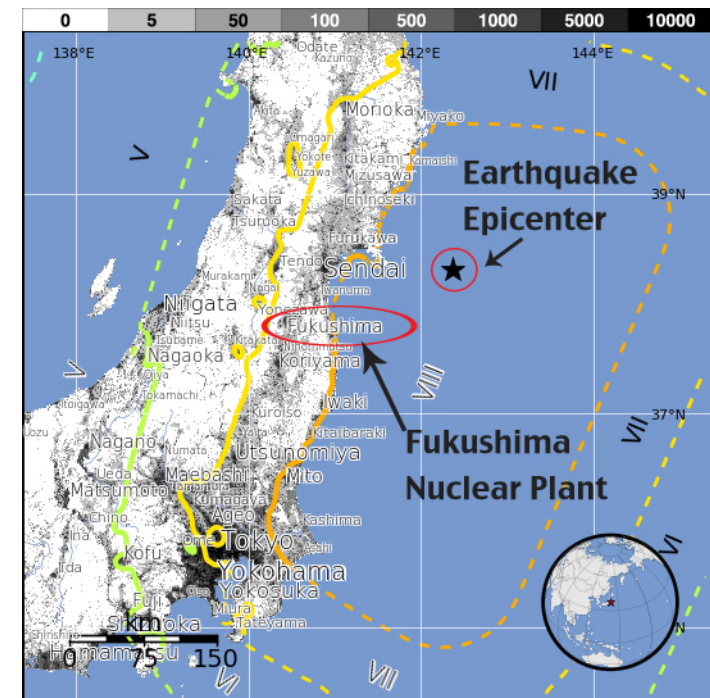
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10 x 10 pixelated neutron detector, 100 ^6Li -glass scintillators ($2 \times 2 \times 9 \text{ mm}^3$)

7 position sensitive 16 channel PMT



Characterization of melted fuel formed in severe nuclear accidents, such as the one at the Fukushima Daiich nuclear power plants



Characterization of melted fuel formed in severe nuclear accidents, such as the one at the Fukushima Daiich nuclear power plants

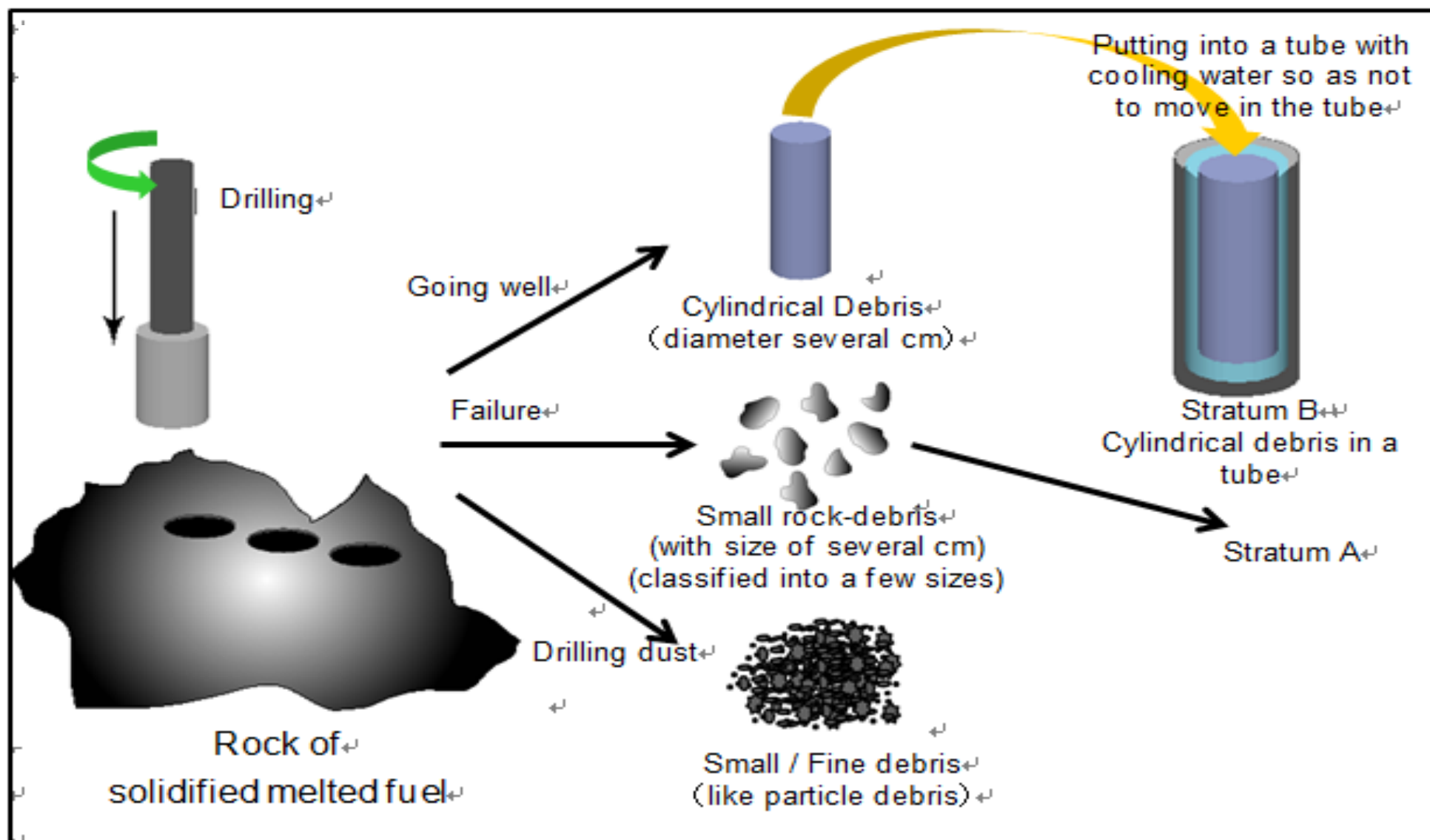
Background

- **Accident at Fukushima following the earthquake on 11/03/2011**
- **Nuclear fuel (NF) will be removed after a cooling period of ~ 10 year**
- **Direct quantification of nuclear materials (NM) in melted fuel (MF) on the site**
- **Quantifying of NM in debris of MF is very complex due to the radioactivity, temperature and variety in size and shape**

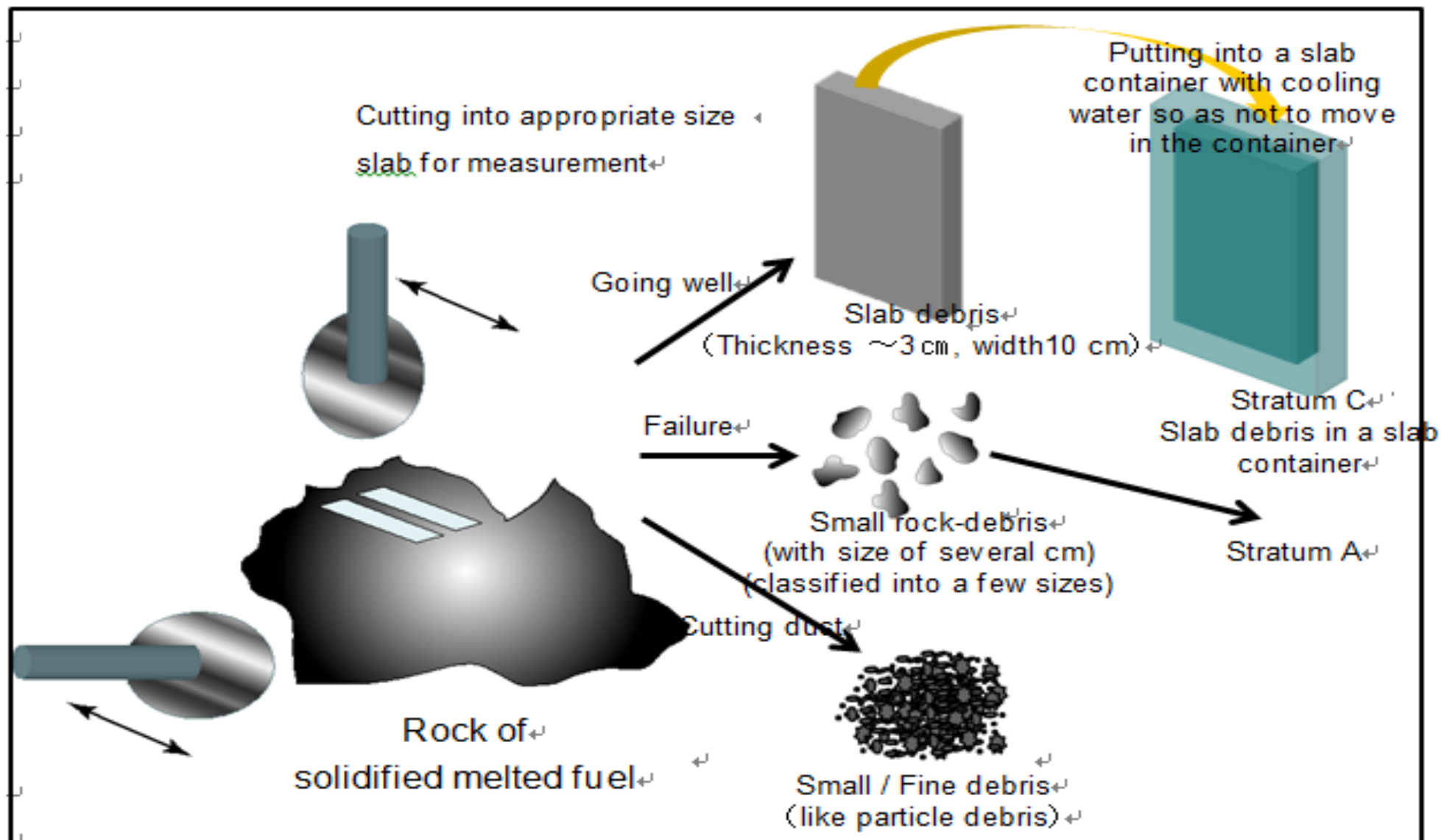
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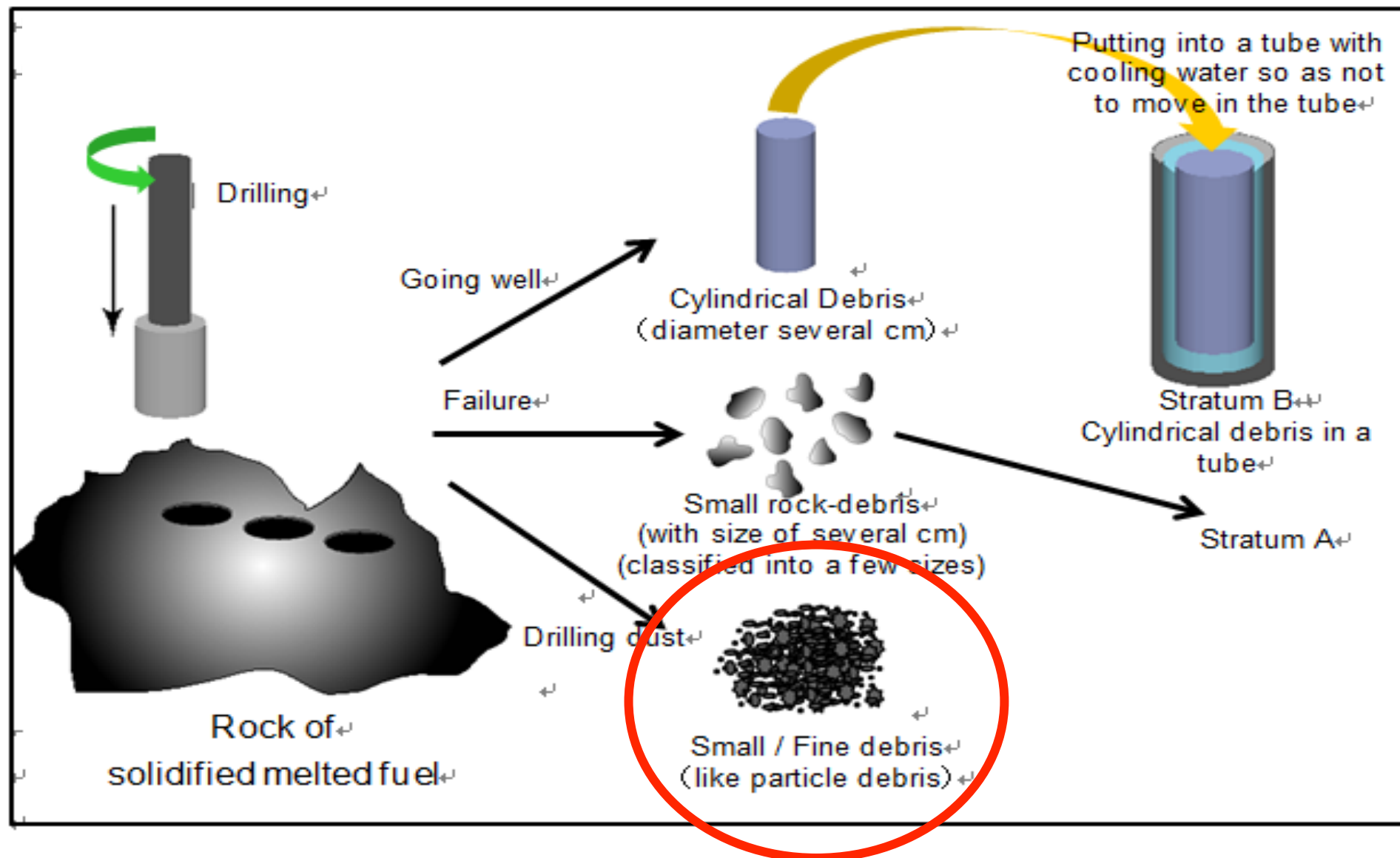
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 - **Direct quantification of nuclear materials (NM) in melted fuel (MF) on the site**
 - **Quantifying of NM in debris of MF is very complex due to the radioactivity, temperature and variety in size and shape**
- ⇒ JAEA proposes to develop Neutron Resonance Densitometry (NRD)**
A method based on NRTA combined with NRCA

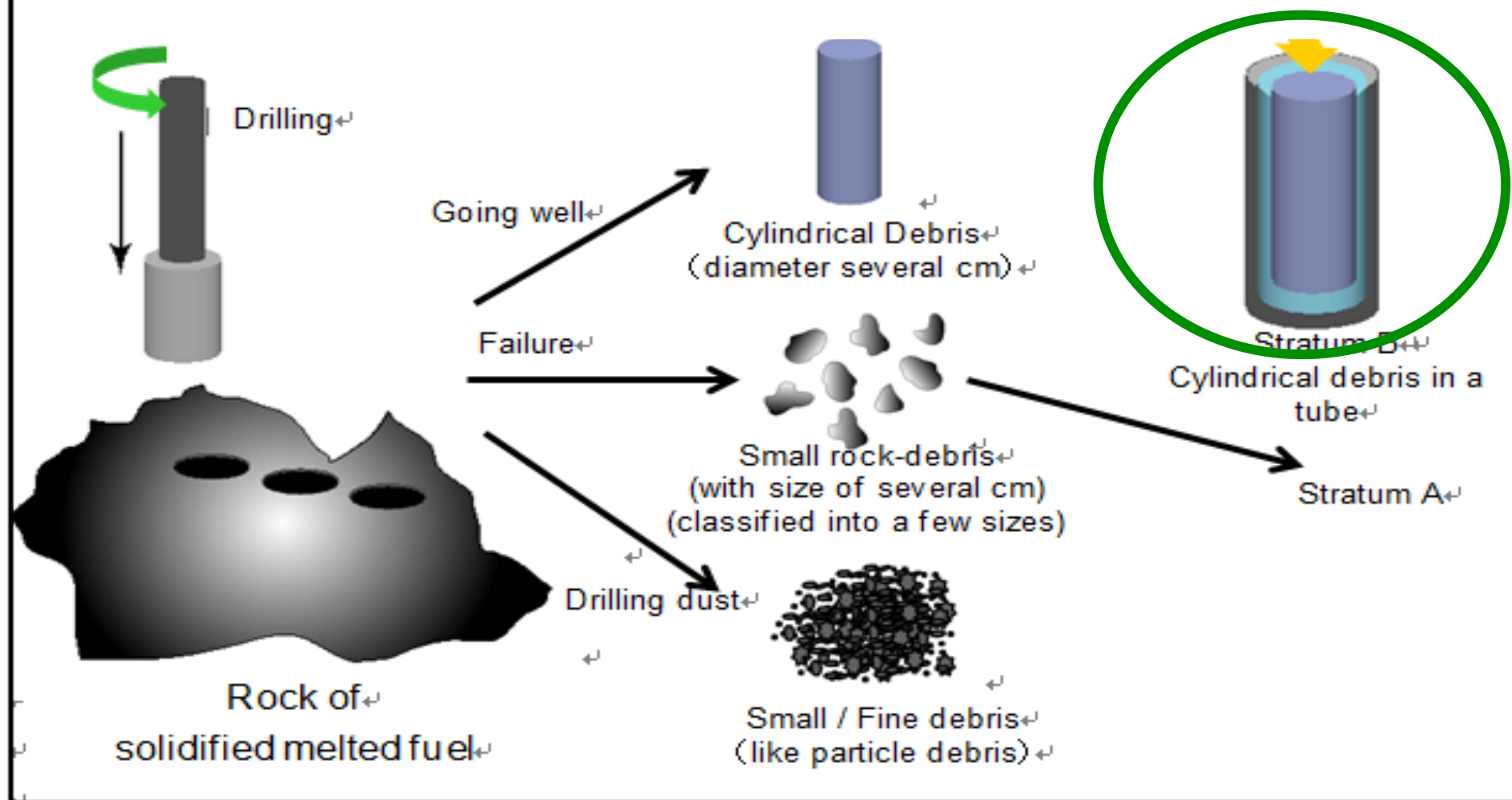


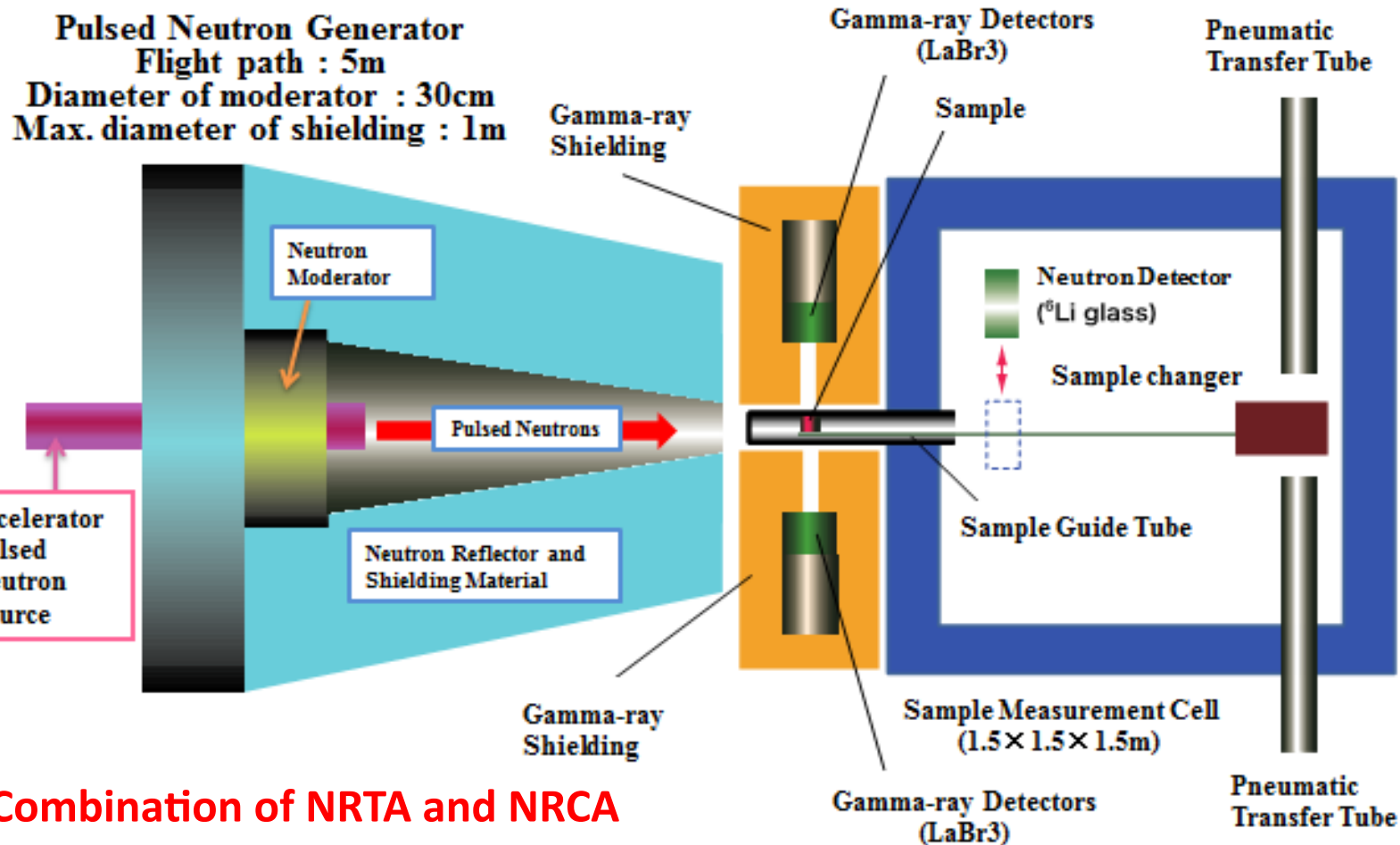
Cutting





Analysis of representative samples

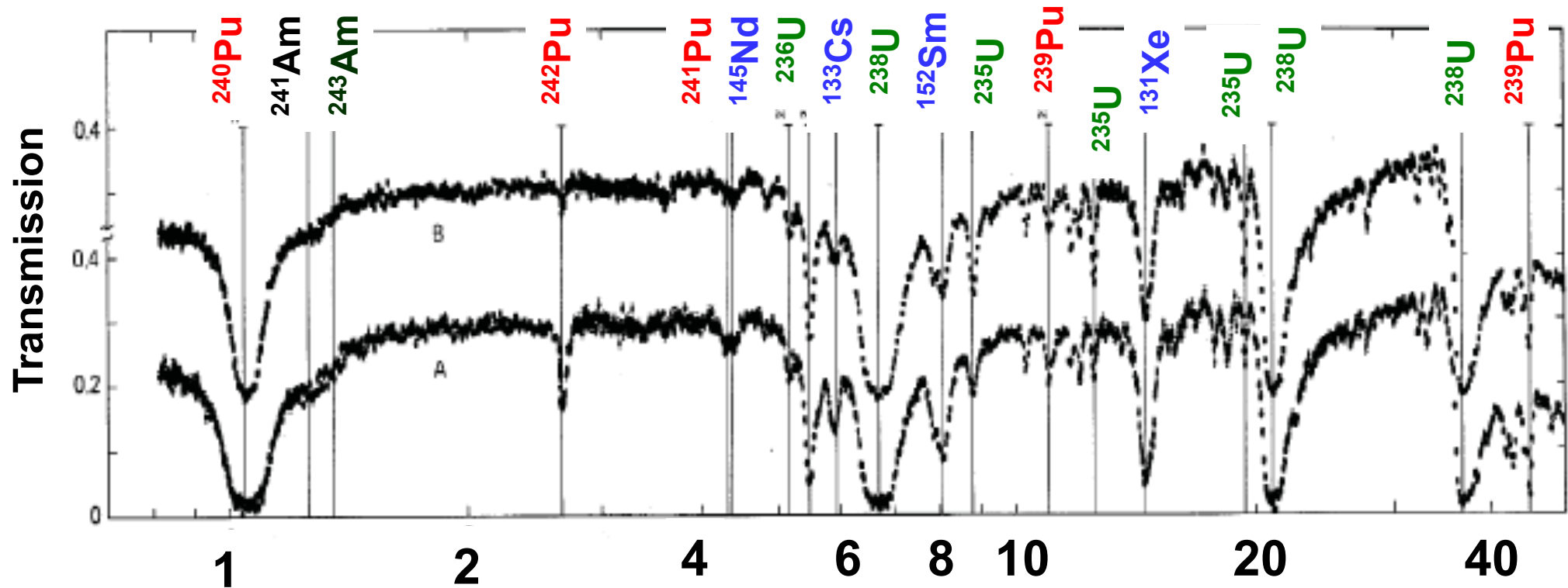




Combination of NRTA and NRCA

**Prototype Neutron Resonance Densitometer
 Designed by JAEA/NSTD**

Use of NRTA to characterize nuclear fuel



A : centre of fuel pin
B : end of fuel pin

■ Study of methodology

- Effect of particle size (• JRC , o JAEA)
- Effect of sample temperature (• JRC , o JAEA)
- Effect of sample thickness (• JAEA , o JRC)
- Effect of contaminated materials (• JAEA , o JRC)
- Study of detection systems (• JAEA , o JRC)
- Generalization of resonance analysis code (REFIT) (• JRC , o JAEA)

■ Study of relevant nuclear data

- Survey of current nuclear data (• JAEA , o JRC)
- Measurement of nuclear data and reference TOF-spectra (• JRC , o JAEA)

■ Study of neutron source

- Electron linear accelerator design (• JAEA , o JRC)
- Neutron source study (• JAEA , o JRC)
- Study of TOF- response function (• JRC , o JAEA)

A. Brusegan[†], F. Corvi, J.C. Drohe, J. Heyse, S. Kopecky, A. Moens, W. Mondelaers, A. Plompen, P. Siegler, D. Vendelbo, R. Wynants and the operators of GELINA
EC - JRC - IRMM (B)

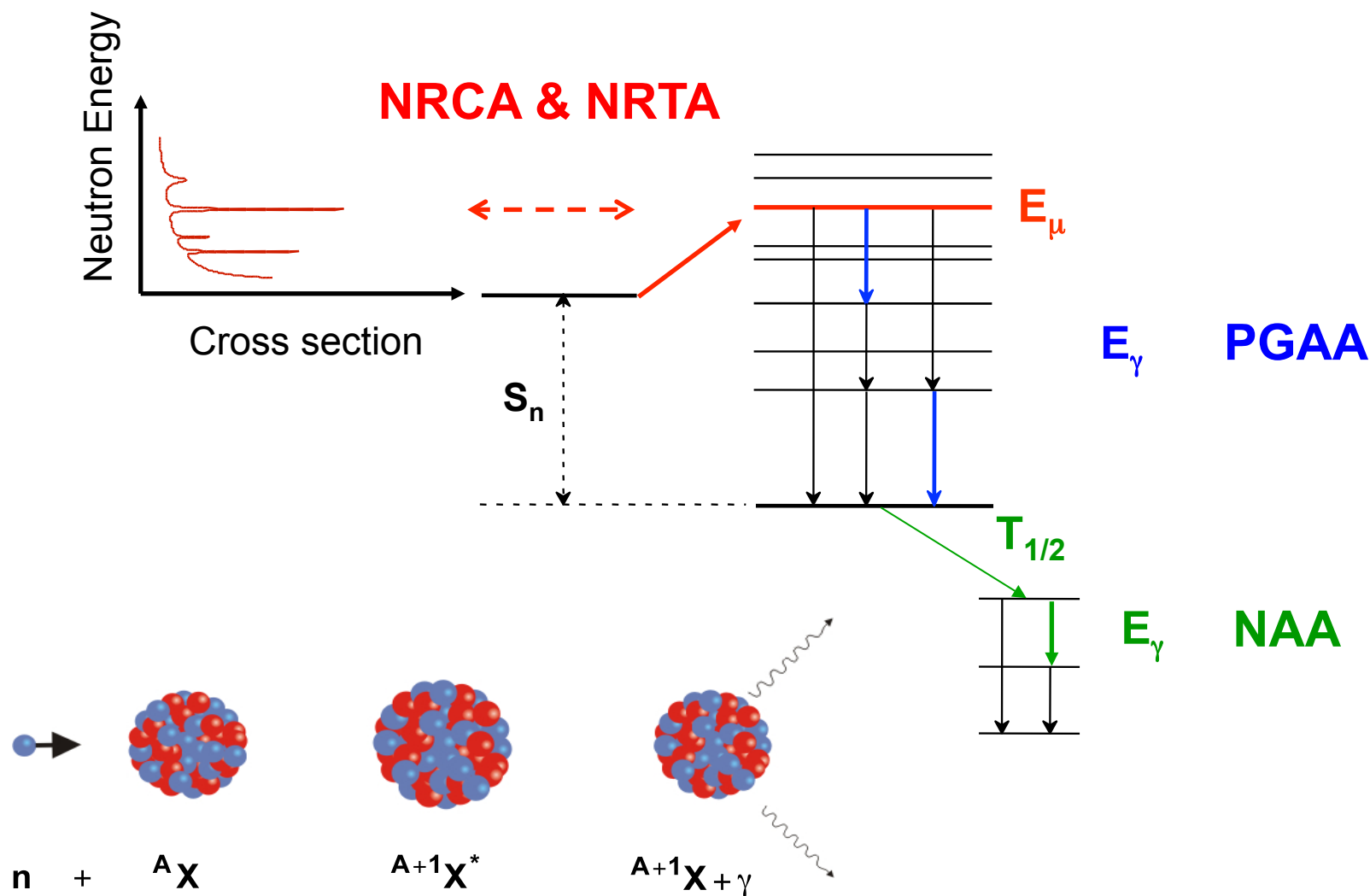
B. Becker, A. Borella, F. Emiliani, K. Kauwenberghs, C. Lampoudis, C. Mihailescu and K. Volev
Current and ex-grantholders, EC - JRC - IRMM (B)

C. Massimi (INFN), M. Moxon, H. Postma and I. Sirakov
Scientific visitors, EC - JRC - IRMM (B)

H. Harada and M. Seya
JAEA (Japan)

Thank you for your attention

Use of resonances as fingerprints



Use of resonances as fingerprints



■ NAA

- Intense thermal neutron flux (irradiation in core, Mol, Delft, NIST, Budapest,...)
- Gamma detector: gamma - ray energy resolution \Rightarrow Ge-detectors

■ PGNAA

- Intense neutron beam, (guided cold neutron beam Budapest, NIST, FRM-II)
- Gamma detector: gamma - ray energy resolution \Rightarrow Ge-detectors

■ NRCA & NRTA

- Pulsed white neutron beam (GELINA, ISIS)
- Gamma/Neutron detector: good time resolution \Rightarrow scintillators

Comparison: NRCA vs. PGAA



■ PGAA

$$k_o = a \beta_\gamma \sigma_\gamma$$

σ_γ **thermal capture cross section**

β_γ **gamma-ray emission probability**

a **isotopic abundance**

$$S_\mu = a A_{\gamma,\mu} \frac{1}{E_\mu}$$

■ NRCA

$A_{\gamma,\mu}$ **capture area**

$$A_{\gamma,\mu} = 4.097 \cdot 10^6 \frac{g\Gamma_n\Gamma_\gamma}{E_\mu\Gamma} (\text{b eV})$$

$1/E_\mu$ **flux shape**

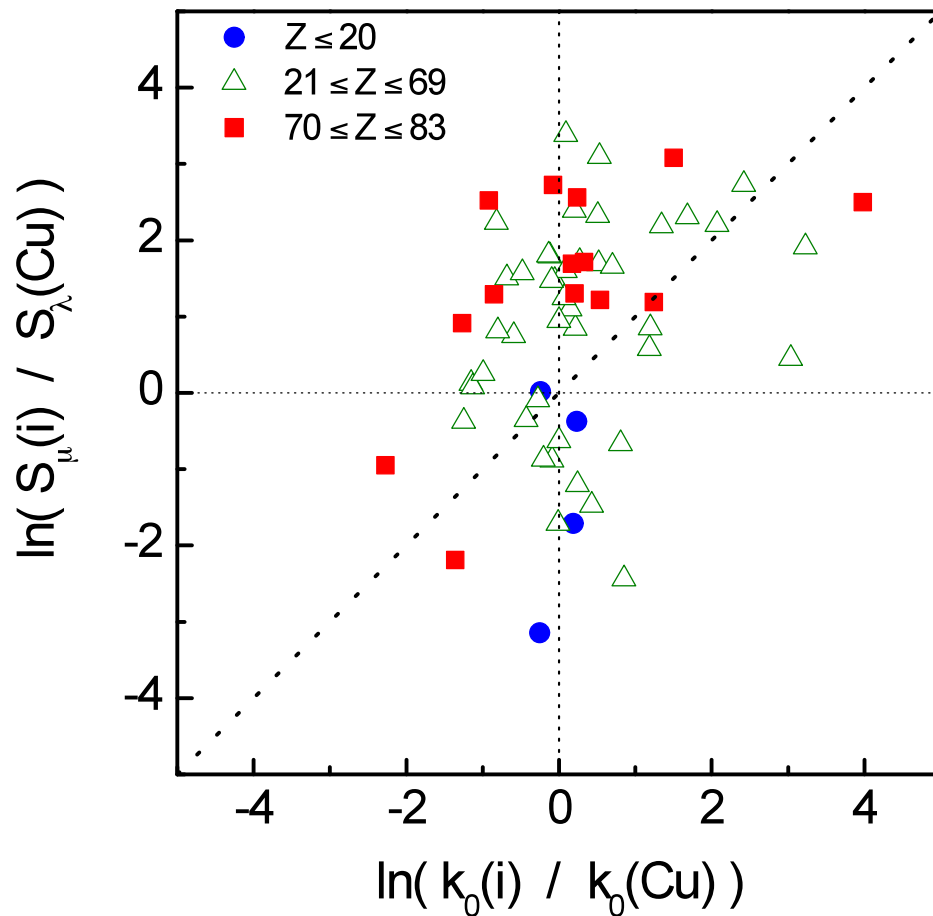
Comparison: NRCA vs. PGAA



PGAA (at Budapest) and NRCA (GELINA)

$\Rightarrow k_0$ and S_μ relative to Cu

Comparable performance for Cu (uncertainty about 1%)



PGAA best for light elements

- H, S, P, and K

NRCA best for heavy elements

- As, Ag, Sb, Au and (Pb)

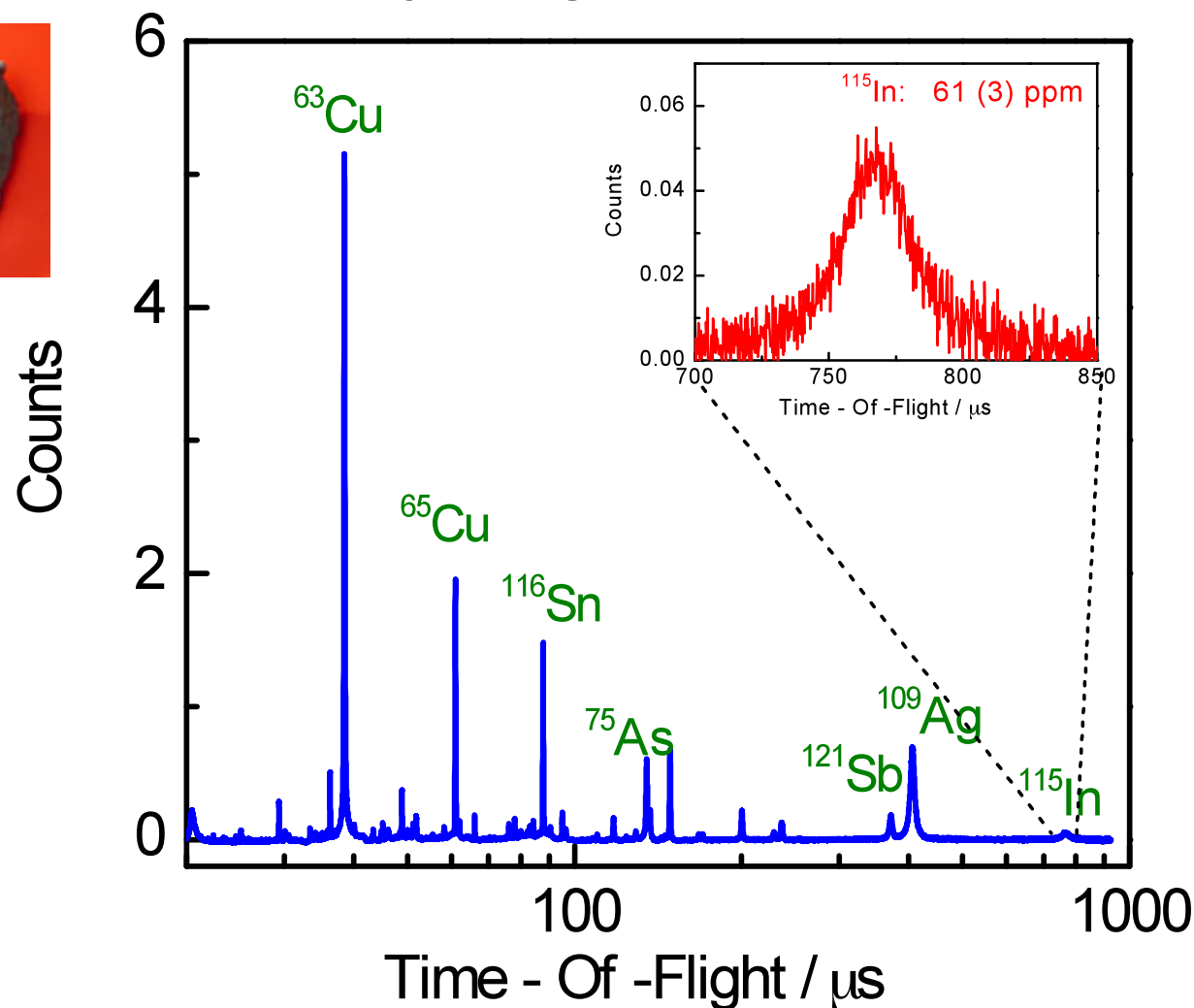
NRTA for light elements + magic

- C, N, O, Na, S and Pb

NRCA : example of TOF-spectrum

Typical dead time : $2.5 \mu\text{s}$

⇒ detection of In is not hindered by strong resonances of other elements (e.g. Cu)



Neutron Production

SHIELDING for
MODERATED SPECTRUM

SHIELDING for
FAST SPECTRUM

