

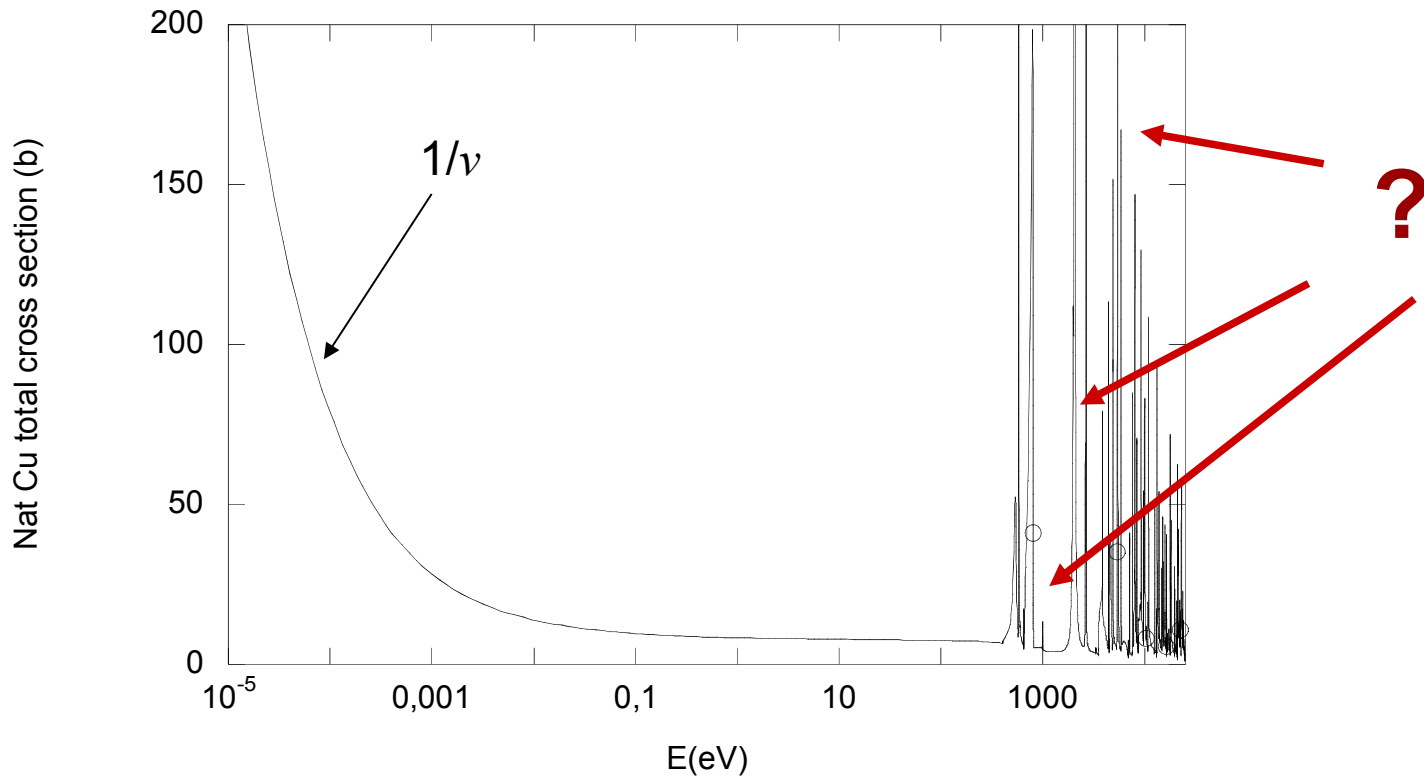


Non-scattering neutron techniques: NRT and NRCA

Enrico Perelli Cippo

“Piero Caldirola” Institut for Plasma Physics – CNR

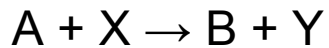
Neutron Resonances: what happens here?



Neutron resonances

better: resonances in neutron absorption and scattering cross-section

Resonance cross-section for the reaction



$$\sigma = \frac{\pi}{k^2} g \frac{\Gamma_{AX} \Gamma_{BY}}{(E-E_R)^2 + \Gamma^2/4}$$

“Breit-Wigner curve”

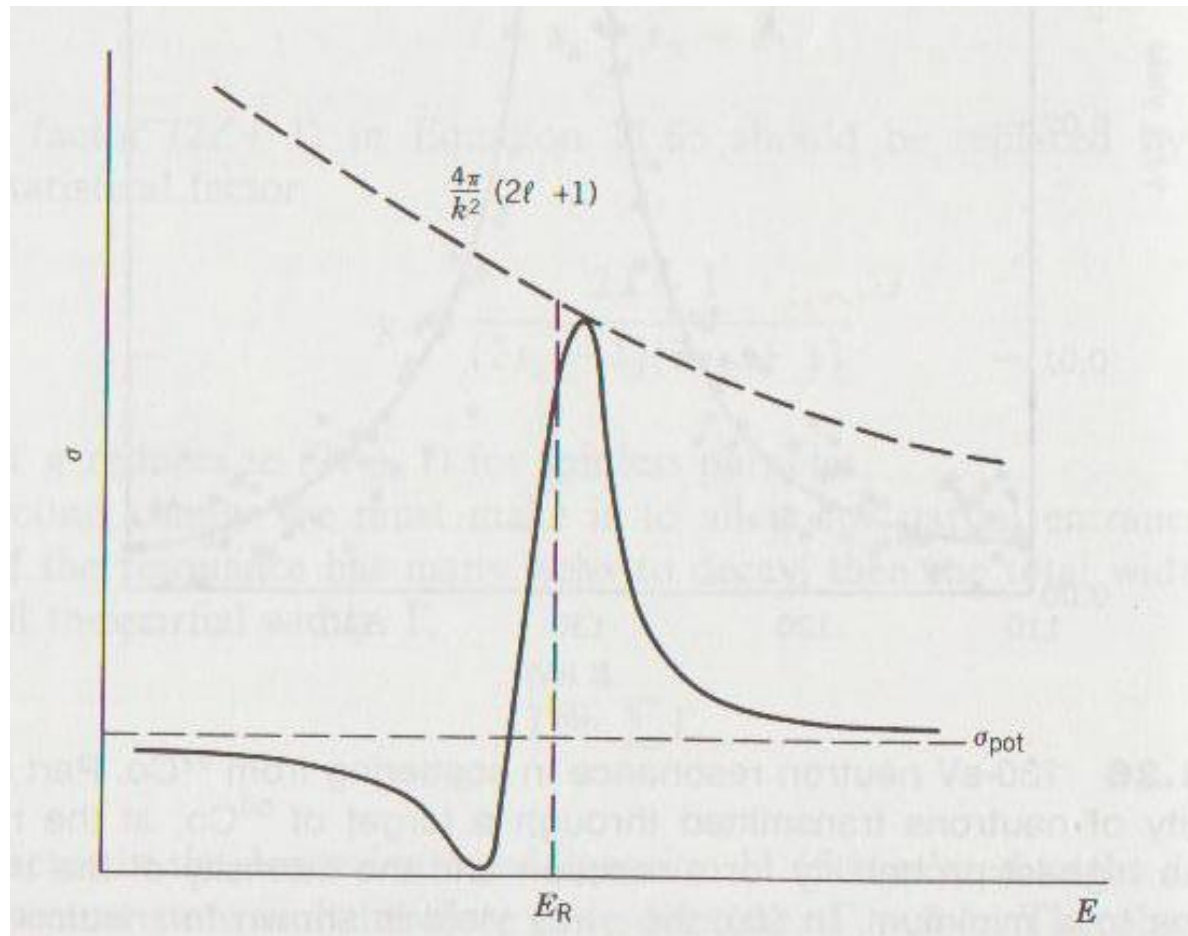
It is related to the existence of a “compound nucleus”.

$$\sigma \sim 1/v$$

off resonance

Neutron resonances

Interference with potential scattering (not related to a compound nucleus) results in slightly asymmetrical total cross section resonances.



Neutron Resonances: who cares about neutron resonances?

Neutron absorption resonances are UNIQUE for every atom and isotope

Neutron absorption resonances can be used for:
Nuclide identification and quantification
Elemental (& isotopic) composition



Neutron Resonance Transmission (NRT)

Neutron absorption resonances can be used for:

Nuclide identification and quantification
Elemental (& isotopic) composition

NRT makes use of epithermal neutrons

E_n up to 1 keV

No sample preparation needed

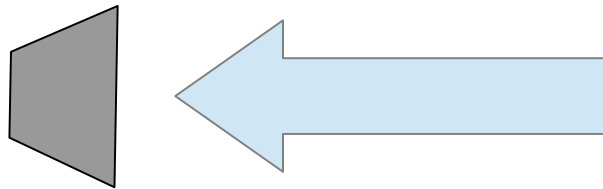
Non - destructive

Negligible residual activation

It can naturally lead to an imaging technique

Neutron Resonance Transmission (NRT)

Neutron detector

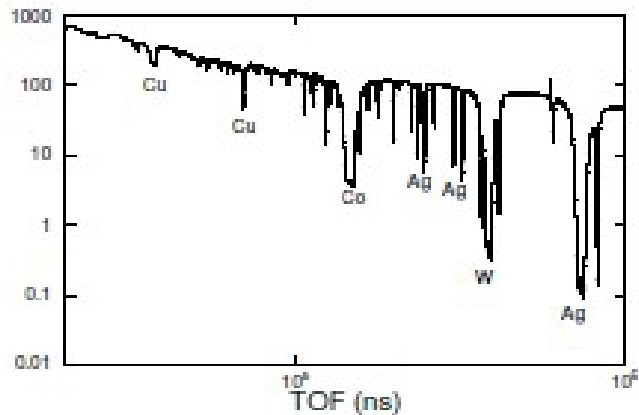


Sample



Neutron beam

TOF DAE



Some previous NRT applications

Investigations on nuclear fuel

Priesmeyer and Harz, “Isotopic content determination in irradiated fuel by neutron transmission analysis” *Atomkernenergie* **25** 109 (1975)

Schrack et al., “Resonance neutron radiography using an electron linac”

IEEE Trans. Nucl. Scie. **28** 1640 (1981)

“

Some previous NRT applications

Investigations on nuclear fuel

Detection of explosives

C. Chen and R.C. Lanza, *IEEE Trans. Nucl. Scie.* **49** 1919 (2002)

Some previous NRT applications

Investigations on nuclear fuel

Detection of explosives

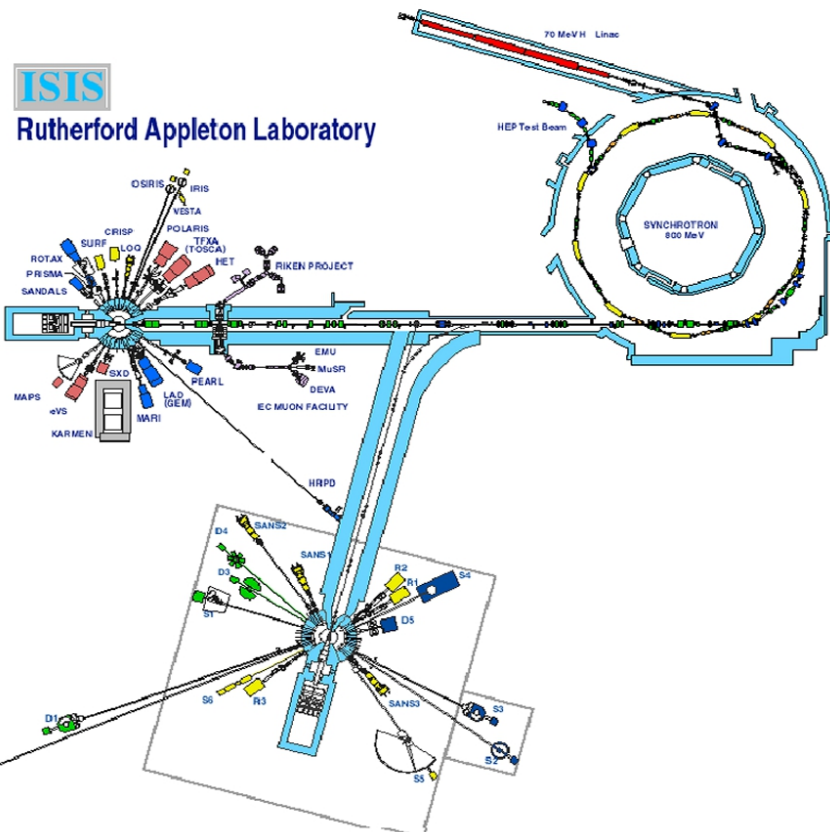
Detection of diamonds in rocks

Watterson and Ambrosi, *Nucl. Instr. Meth. A* **513** 367 (2003)

NRT makes use of epithermal neutrons

E_n up to 1 keV

Pulsed white neutron beam (GELINA, ISIS)

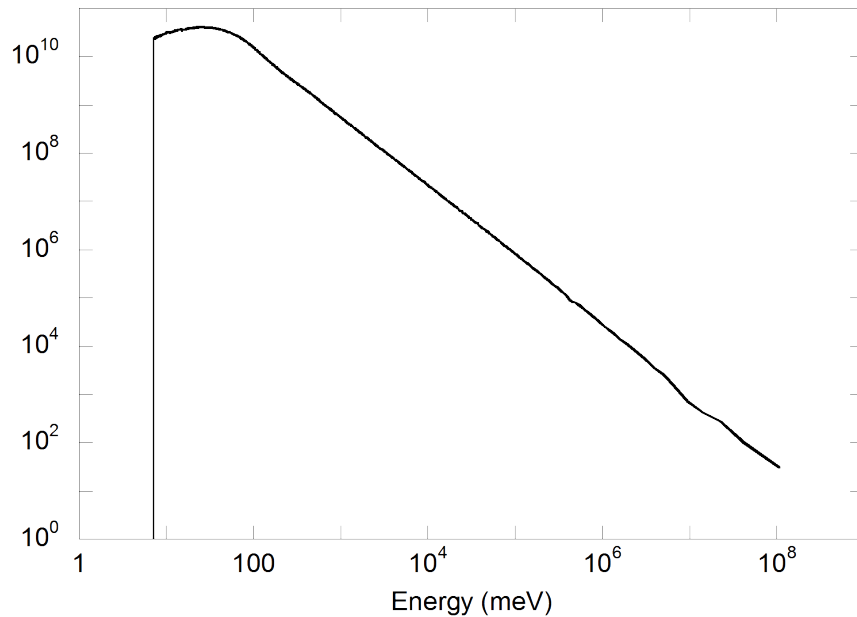


INES beamline at the ISIS spallation source	
Average current	150 – 180 μA (p)
Neutron pulse width	300 ns
Flight path length	23.0 m
Beam dimensions	50 x 50 mm
Flux at sample pos.	10^3 n/eV s cm ² at 10 eV

NRT makes use of epithermal neutrons

E_n up to 1 keV

Pulsed white neutron beam (GELINA, ISIS)



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“Undermoderated” beam at
INES

NRT makes use of epithermal neutrons

E_n up to 1 keV

Pulsed white neutron beam (GELINA, ISIS)

Absorption resonances are identified with the TOF technique

$$E_n = \left(\frac{72.2985L}{t + t_0} \right)^2$$

FAST detectors and electronics are crucial

And if you want images, you also must consider spatial resolution

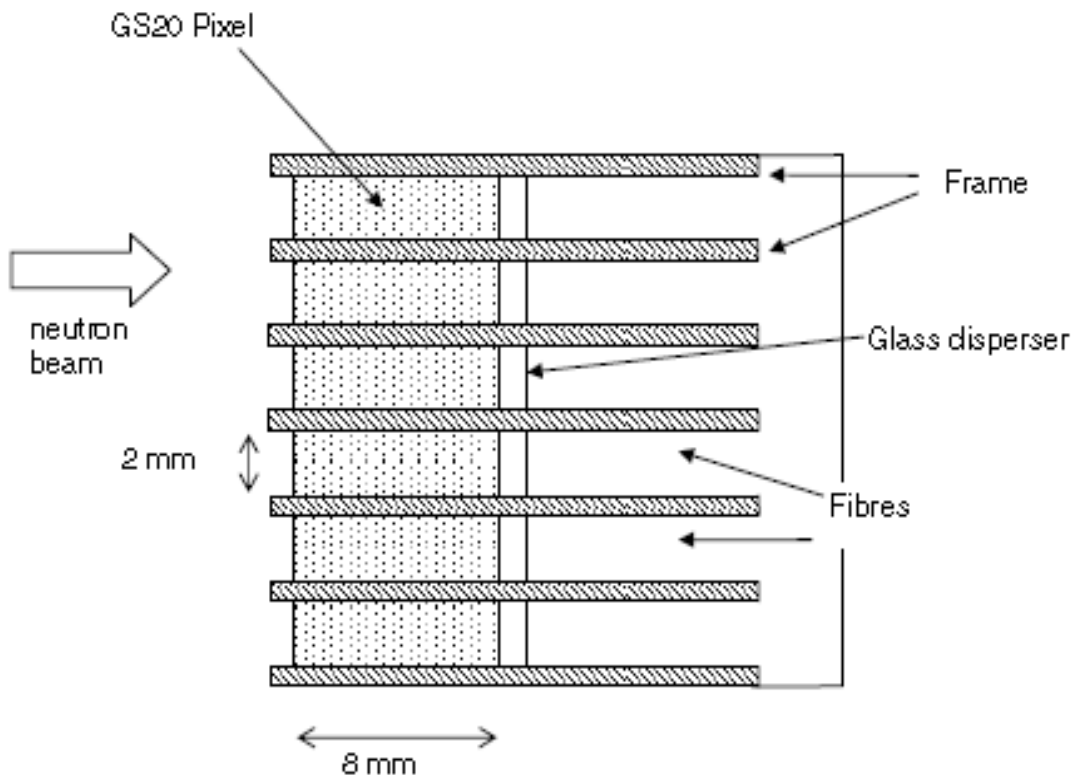
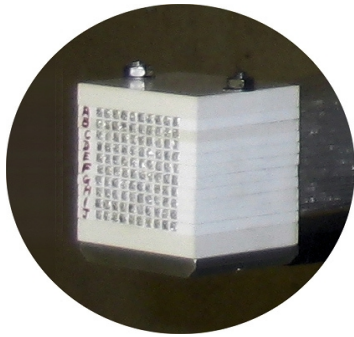
PSND

10 x 10 *GS20* Li-enriched scintillator pixels

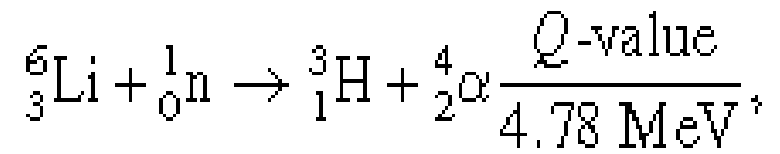
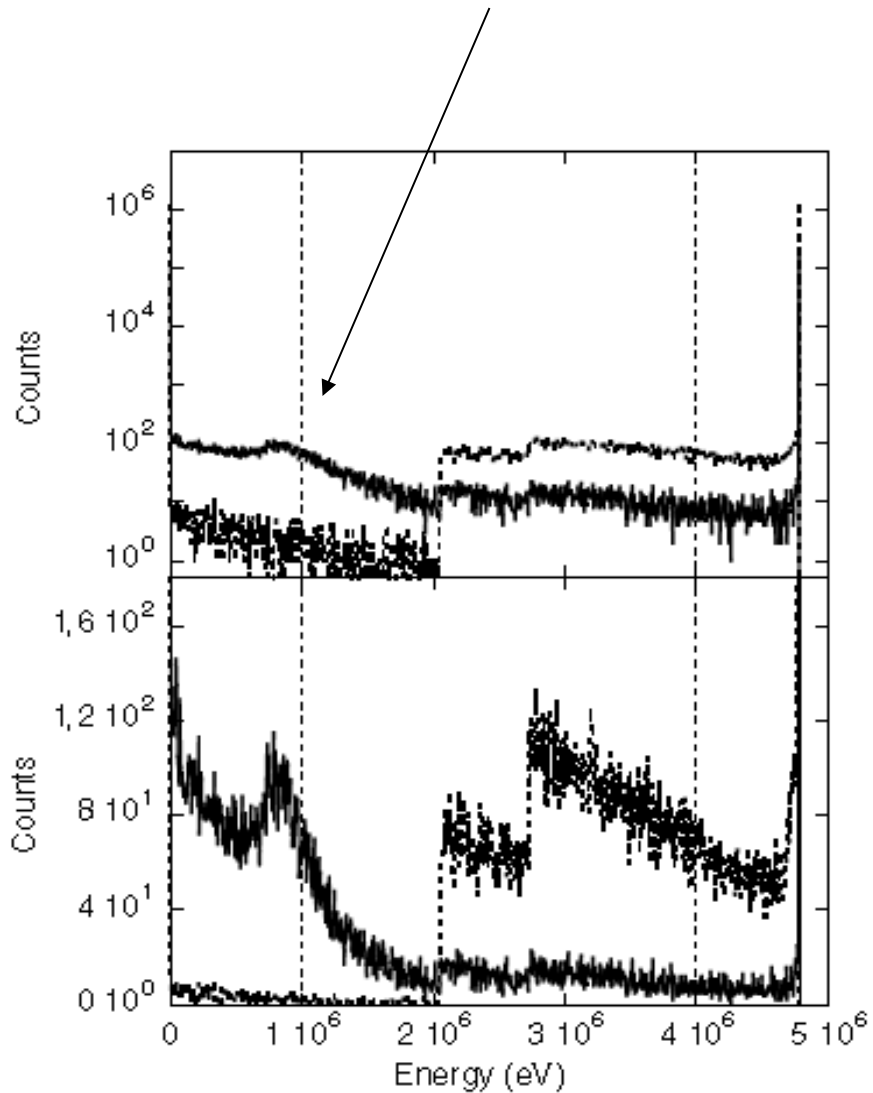
25% efficiency at 10 eV

4 % efficiency at 1 keV

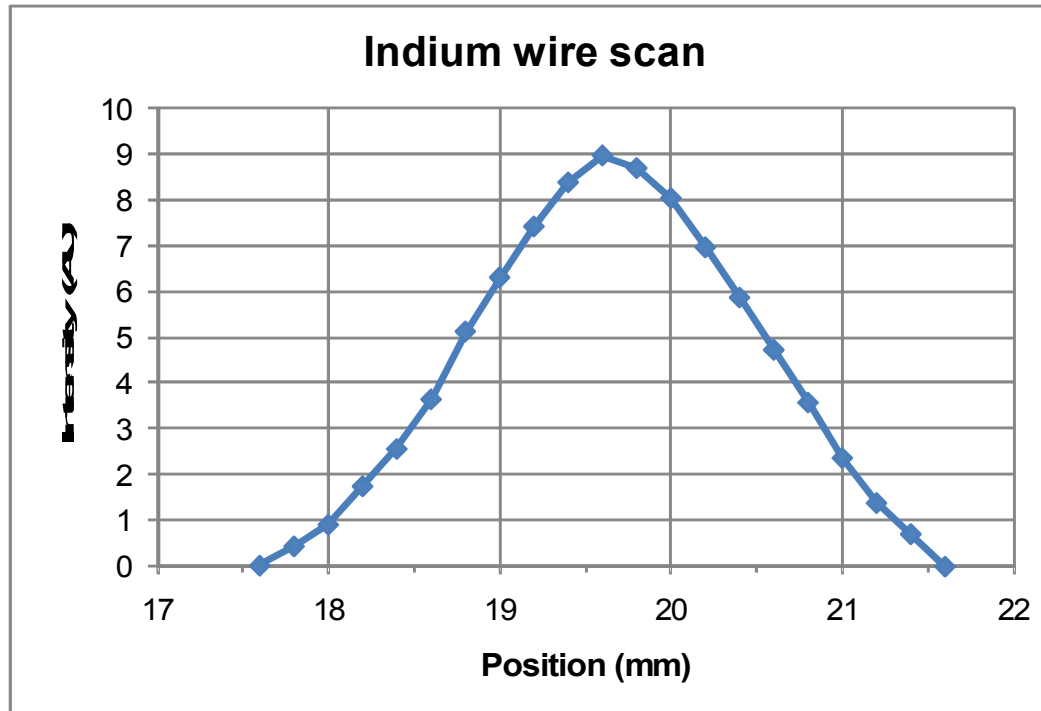
FAST response



“Inverted wall effect” causes
cross-talk between pixels

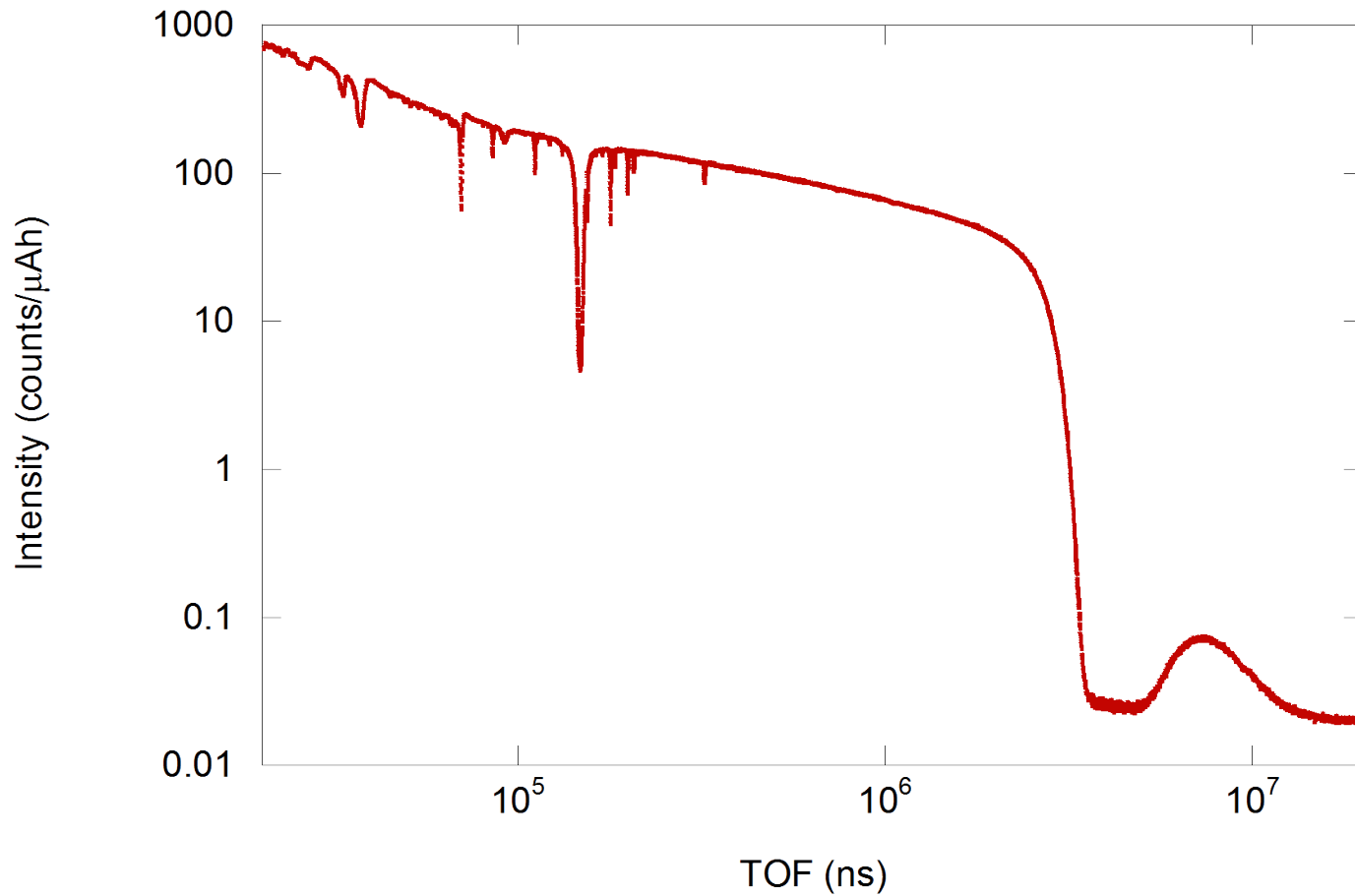


Spatial resolution

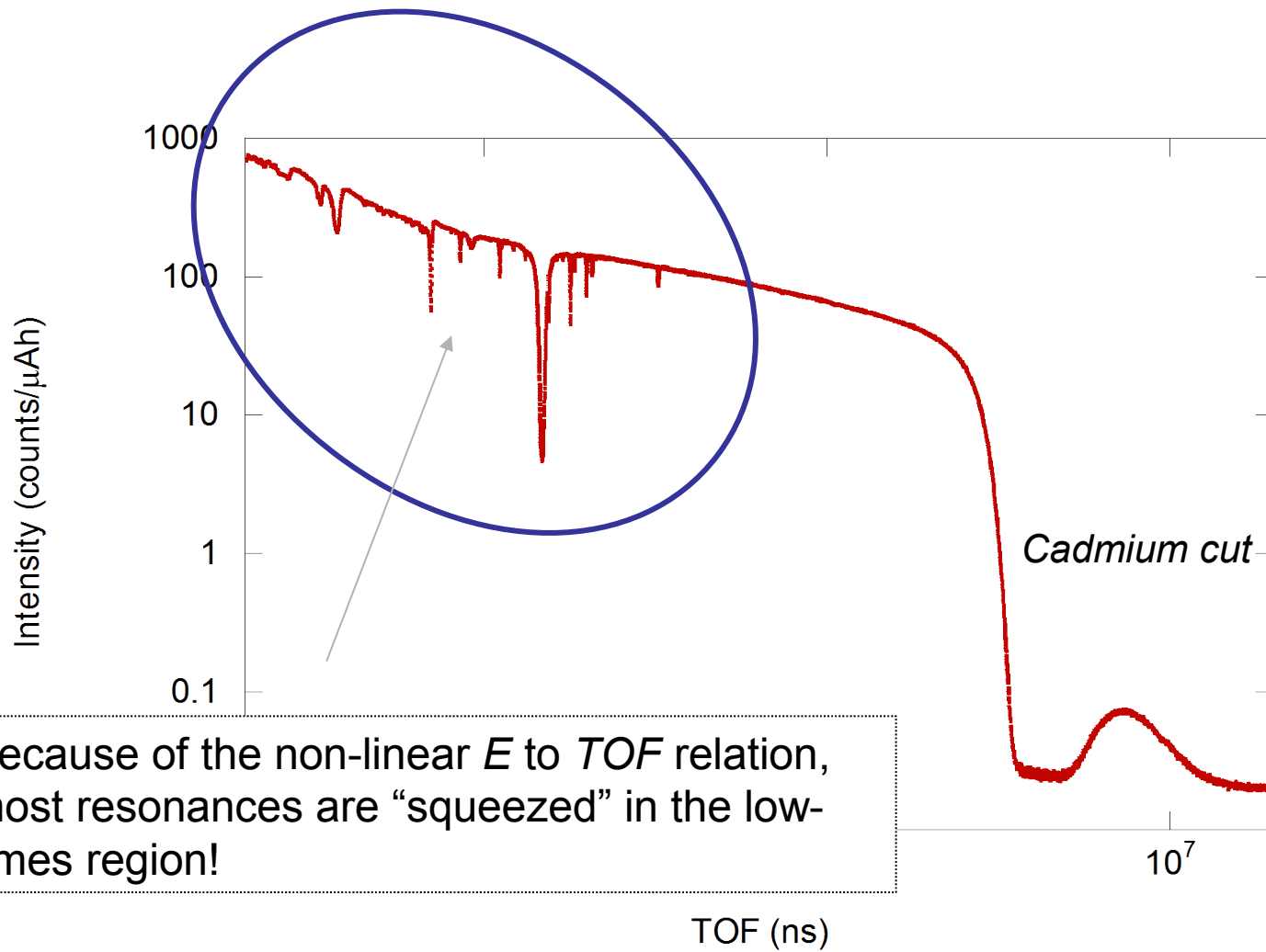


Space resolution of about 2.5 mm FWHM

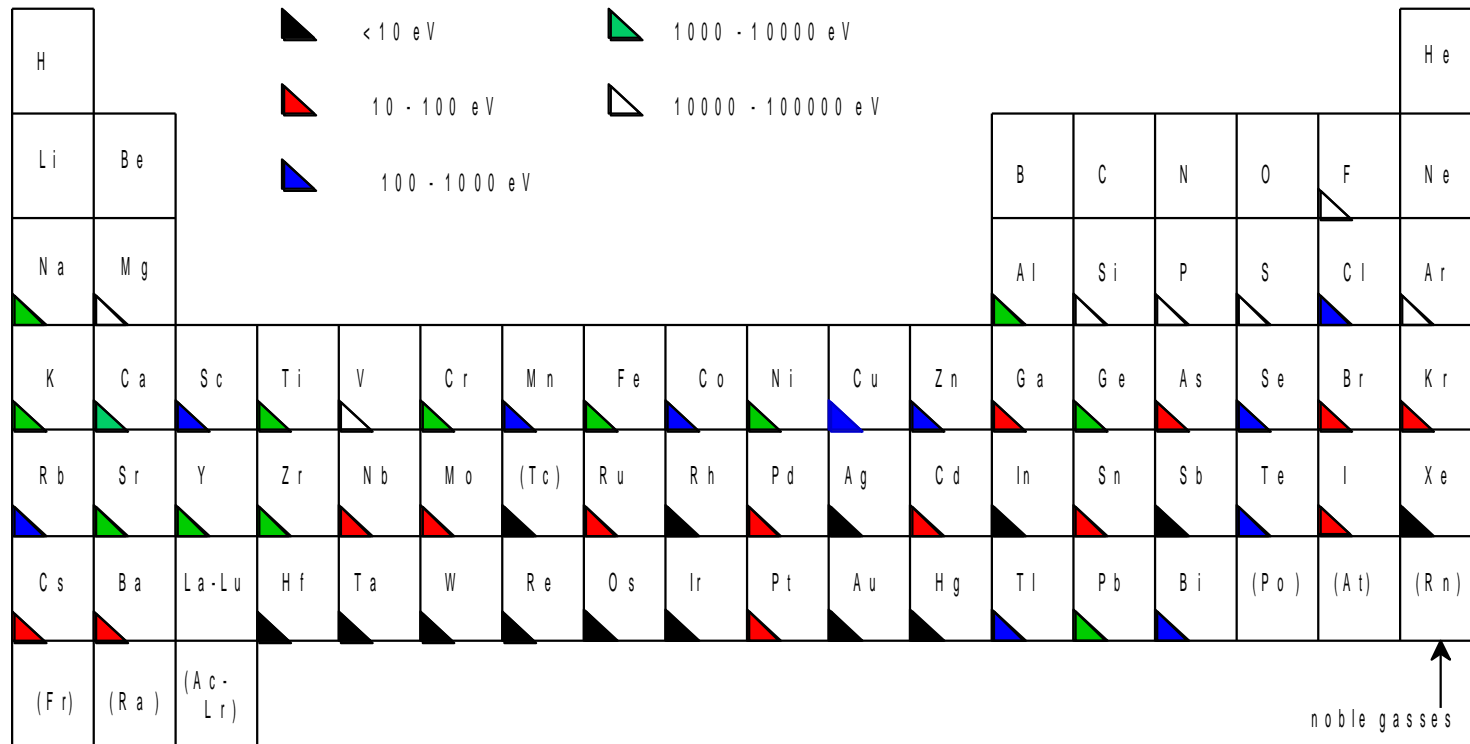
The transmission TOF spectrum



The transmission TOF spectrum



Many elements of CH interest have resonances in the slightly lower energy (slightly longer TOF....) region.....



Lanthanides
(Rare Earth elements)

La	Ce	Pr	Nd	(Pm)	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb	Lu
(Ac)	Th	(Pa)	U	(Np)	(Pu)									

Actinides

The *optimum* NRT analysis approach

That sounds ironic...



The physical quantity measured is the transmission factor:

$$T(E) = \frac{C_{in}}{C_{out}} = e^{-\sum_x n_x \sigma_{tot}(E)} R(E)$$

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$$T(E) = \frac{C_{in}}{C_{out}} = e^{-\sum_X n_x \sigma_{tot}(E)} R(E)$$

$$T_{exp}(t) = N \frac{S_{in}(t) - B_{in}(t)}{S_{out}(t) - B_{out}(t)}$$

where $S_{in, out}$ are the signal obtained from the sample-in and sample-out for the flux, and $B_{in, out}$ are the background levels in the two cases. N is a normalisation constant.

This is valid for all the values of t (in and off-resonances).

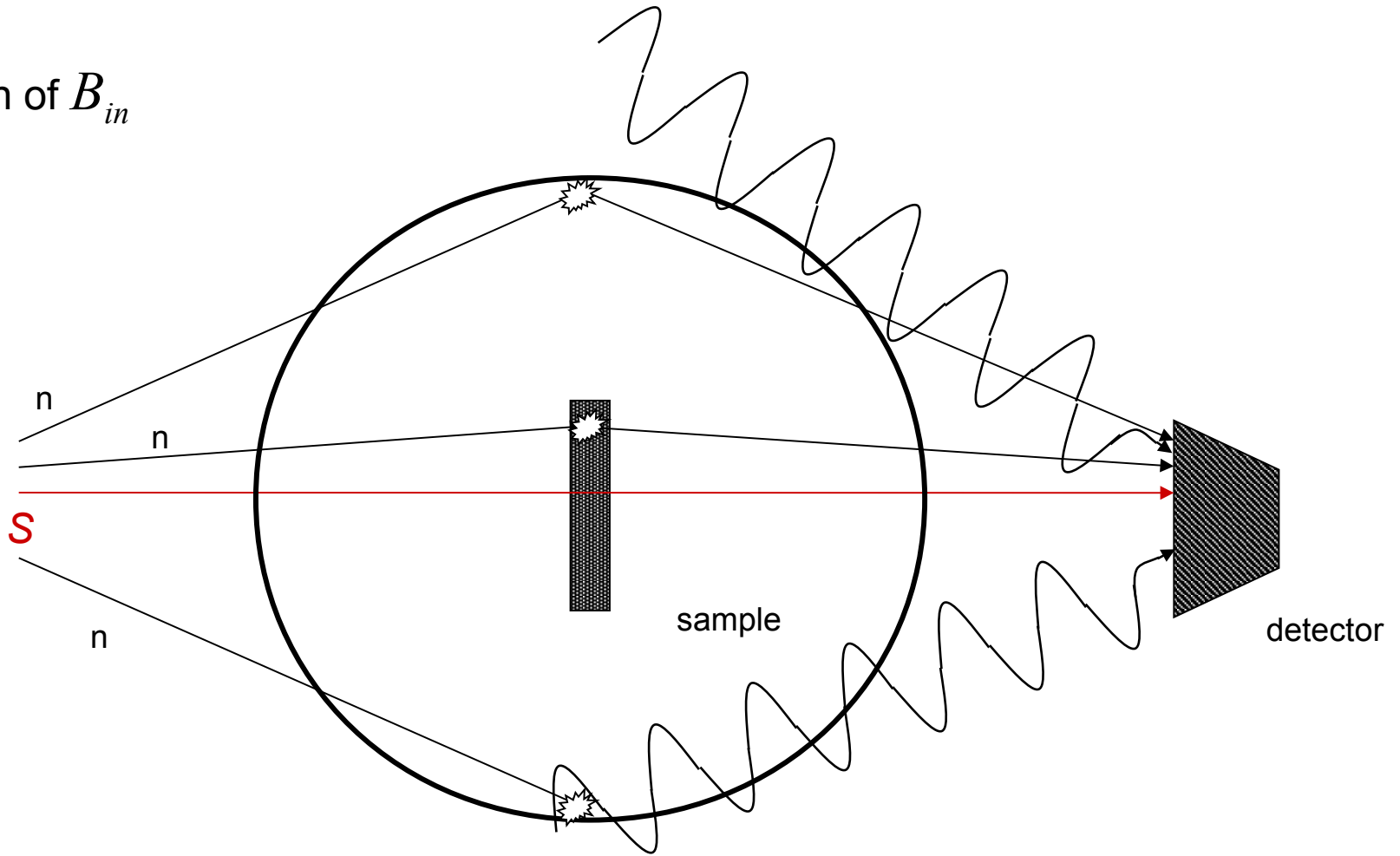
The physical quantity measured is the transmission factor:

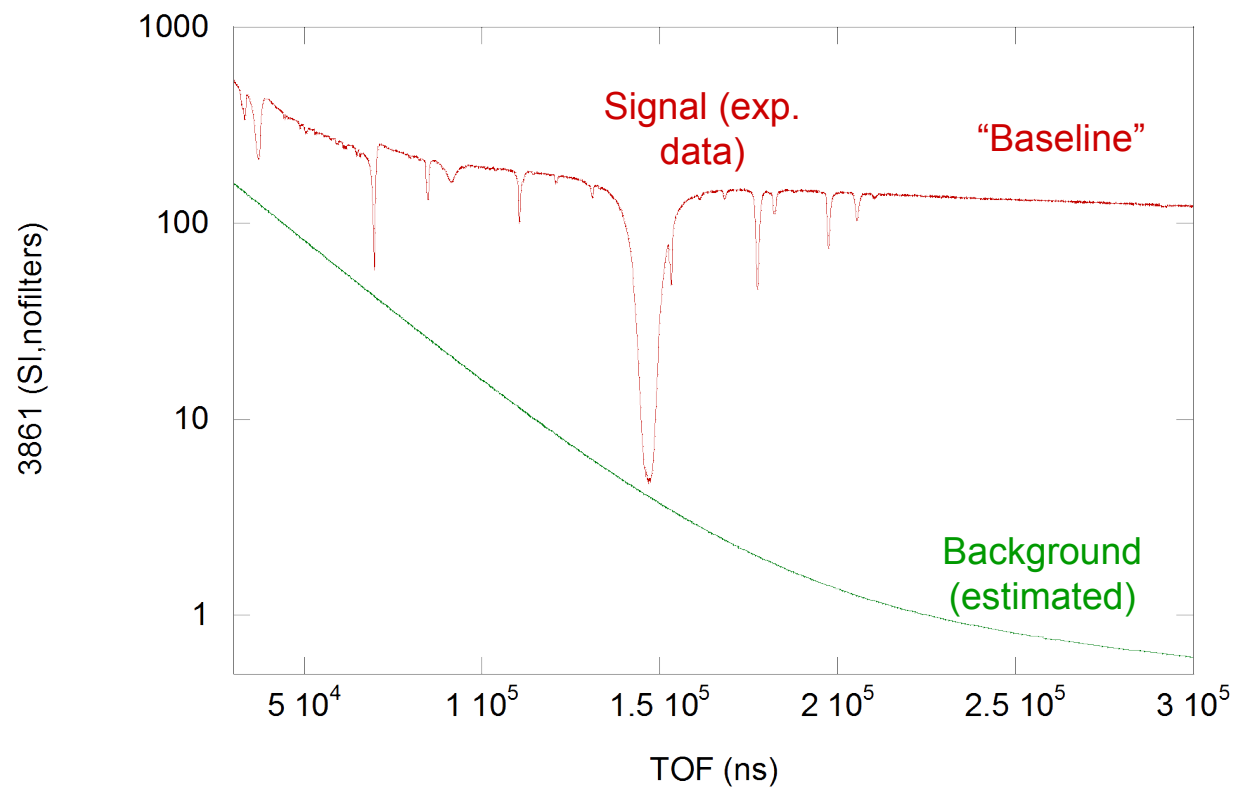
$$T_{\text{exp}}(t) = N \frac{S_{\text{in}}(t) - B_{\text{in}}(t)}{S_{\text{out}}(t) - B_{\text{out}}(t)}$$

experimental data

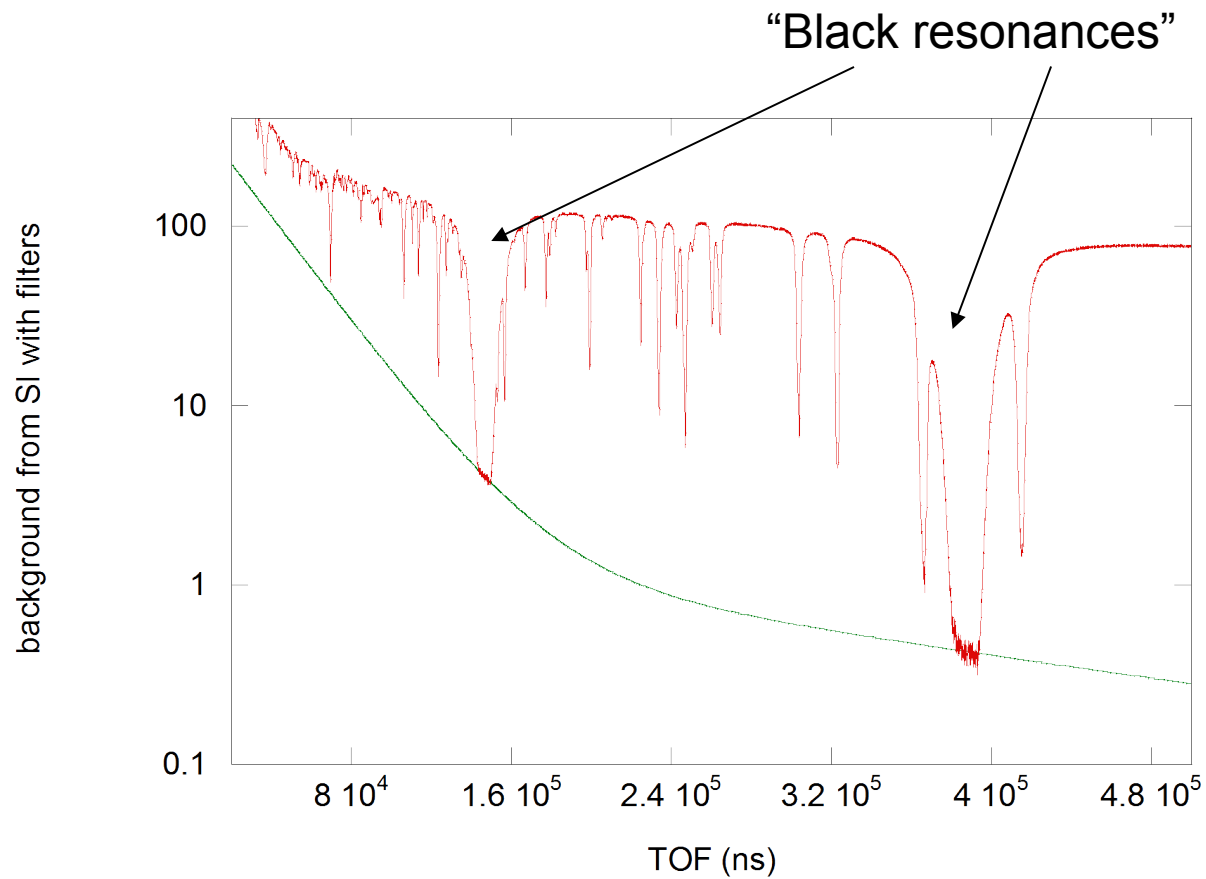
evaluated analytical function

Definition of B_{in}



Definition of B_{in} 

How to determine B_{in}



How to determine B_{in}

Resonances used:

Bi 800 eV 59 μ s

Co 132 eV 150 μ s

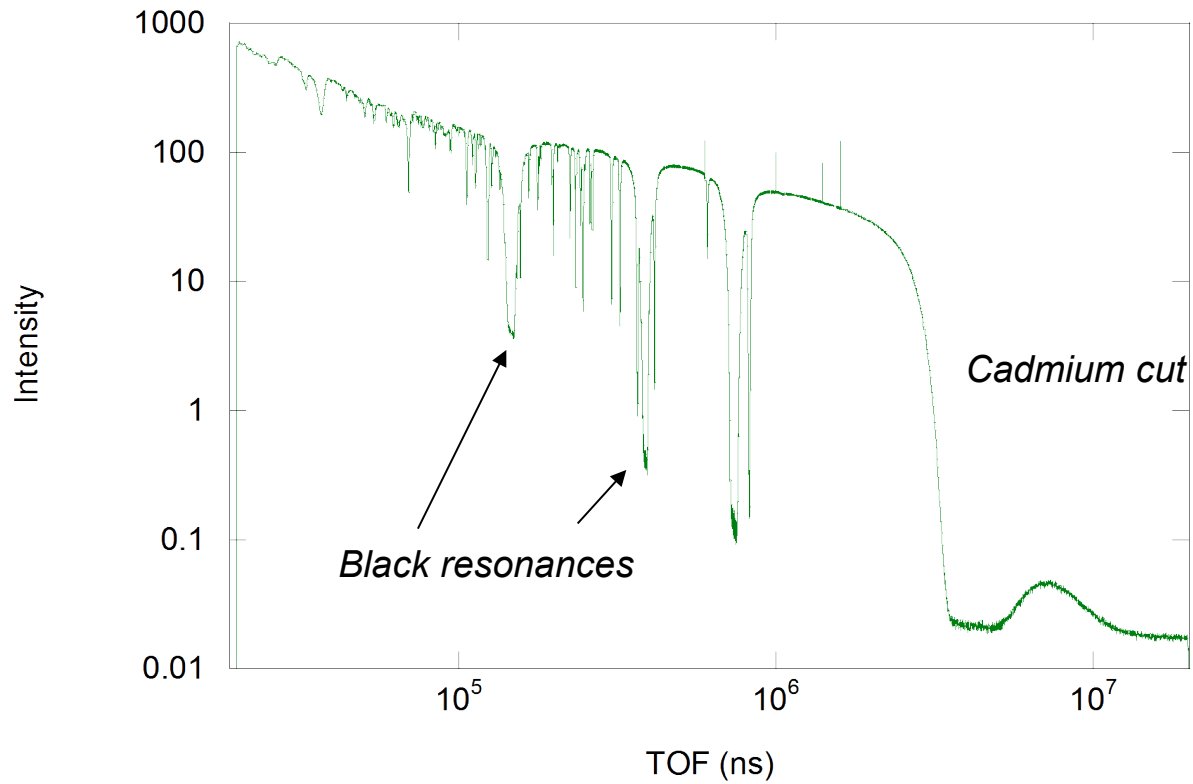
W 19 eV 390 μ s

Ag 5 eV 750 μ s

Cd 0.5 eV 3000 μ s (also for reducing activation by thermal neutrons)

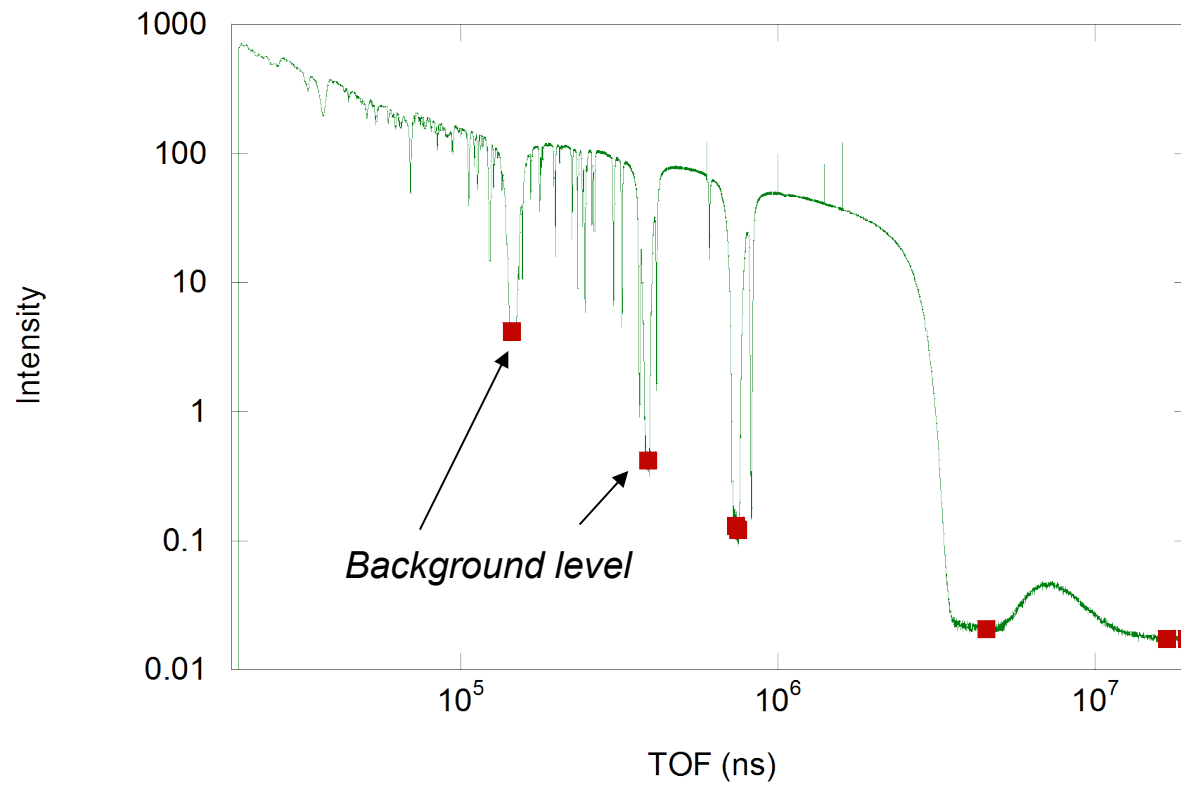
How to determine B_{in}

Spectrum from a sample-in + black resonances run



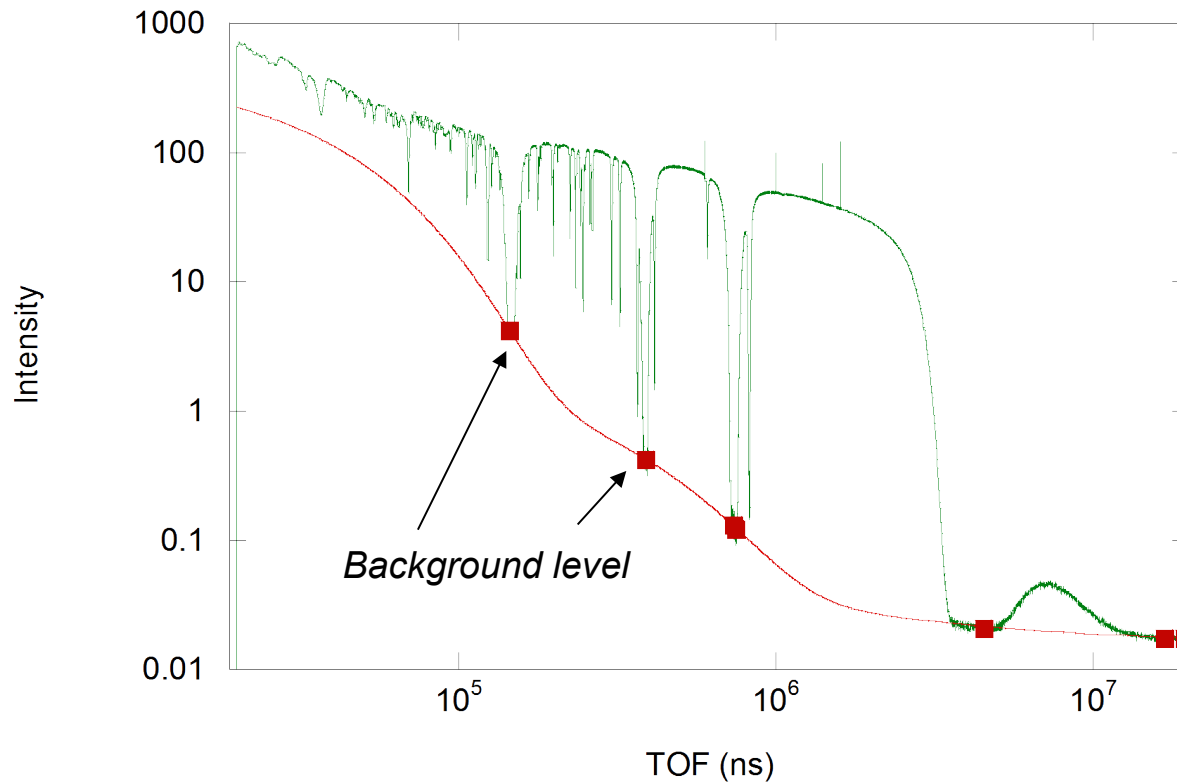
How to determine B_{in}

Estimated background level



How to determine B_{in}

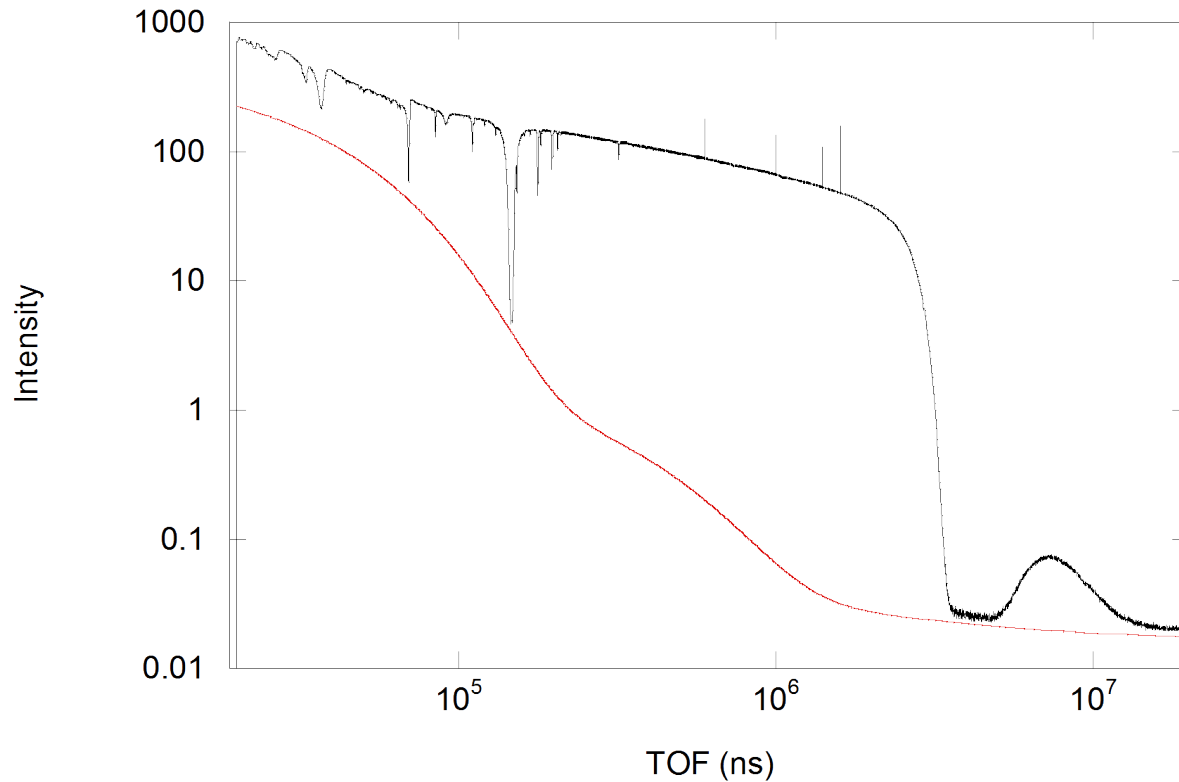
Estimated background interpolation



$$a + b \exp(-ct) + d \exp(-et) + f t^g$$

How to determine B_{in}

Estimated background level B_{in} compared with Sample-in (no filters) S_{in}



S_{in} is the TOF transmission spectrum of the sample that has undergone

- Dead time correction:

$$DTCF = \frac{C_o}{(1 - \beta_o)} ; \quad \beta_o = \sum_{i=(2t-1)/2}^{((2t-1)/2)+\Delta t} N(t)$$

$$\Delta t \approx 275 \text{ ns}$$

DT correction function is an integral function in t



M. S. Moore, "Rate dependence of counting losses in neutron time-of-flight measurements", *Nucl. Instr. Meth.* **169** 245 (1980)

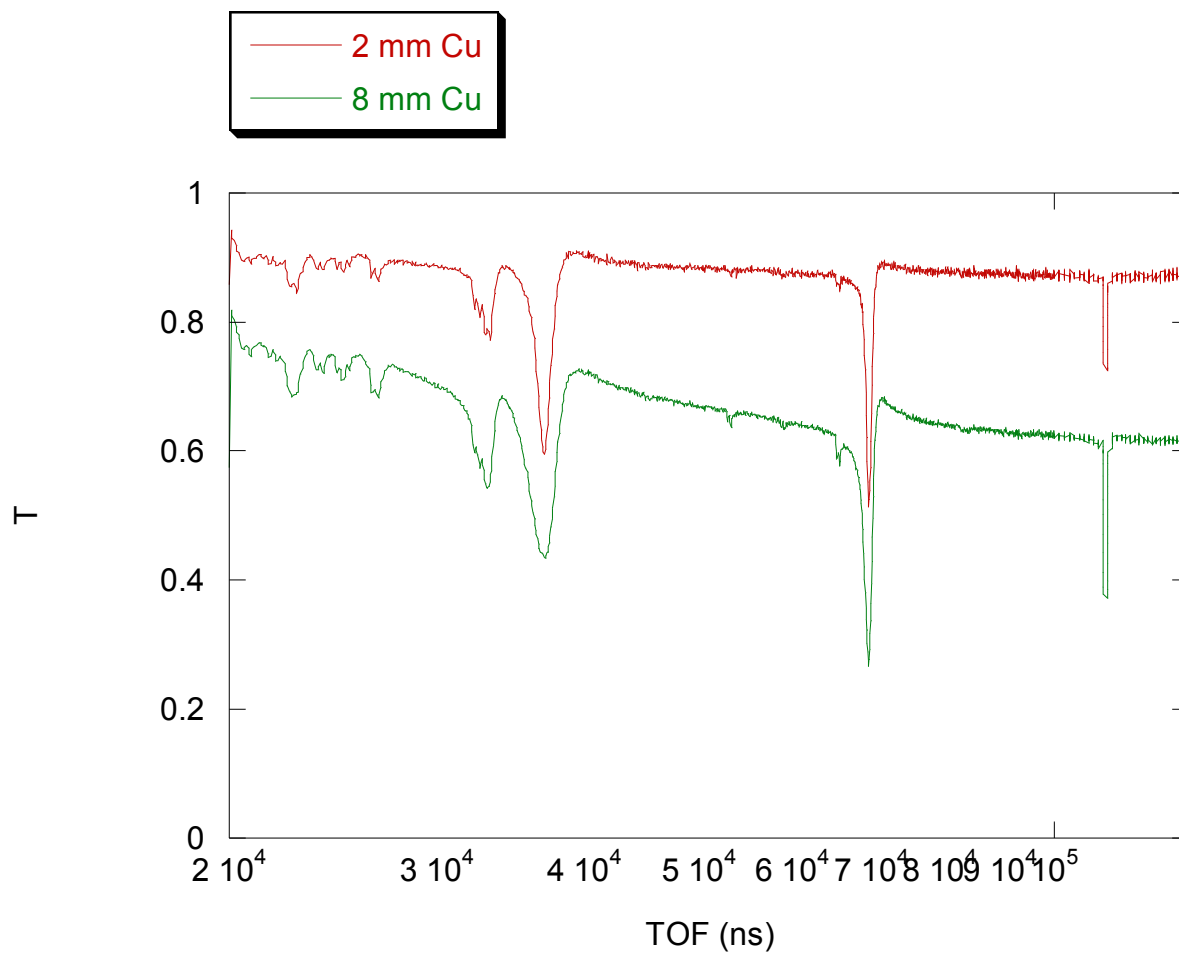
S_{in} is the TOF transmission spectrum of the sample that has undergone

- Dead time correction
- Normalisation to neutron flux

S_{in} is the TOF transmission spectrum of the sample that has undergone

- Dead time correction
- Normalisation to neutron flux
- **Normalisation to bin width**

Transmission factor for Cu samples (INES, may – june 2009)



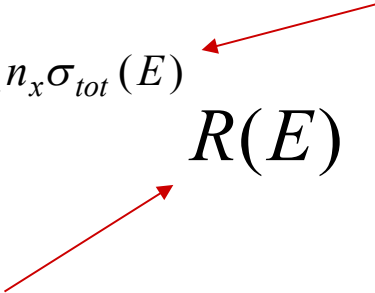
Optimum NRT analysis approach: REFIT

REFIT is a toolkit for analysis of transmission (and capture) neutron resonances spectra.

Advantage: fully quantitative

$$T(E) = \frac{C_{in}}{C_{out}} = e^{-\sum_x n_x \sigma_{tot}(E)} R(E)$$

Resonances are Doppler broadened



Data do not have infinitely good resolution

Optimum NRT analysis approach: REFIT

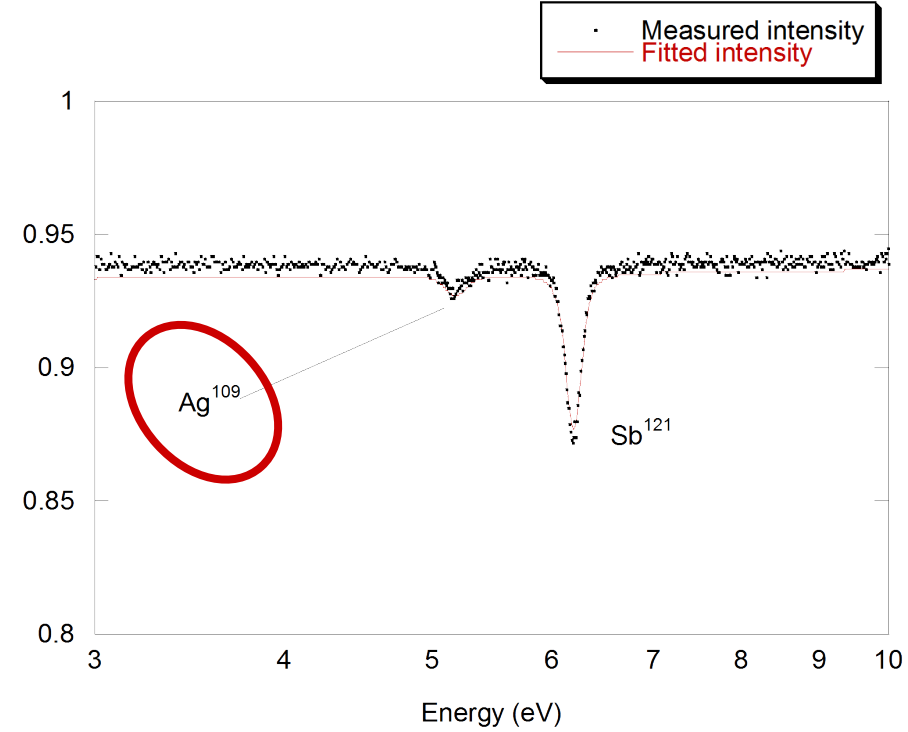
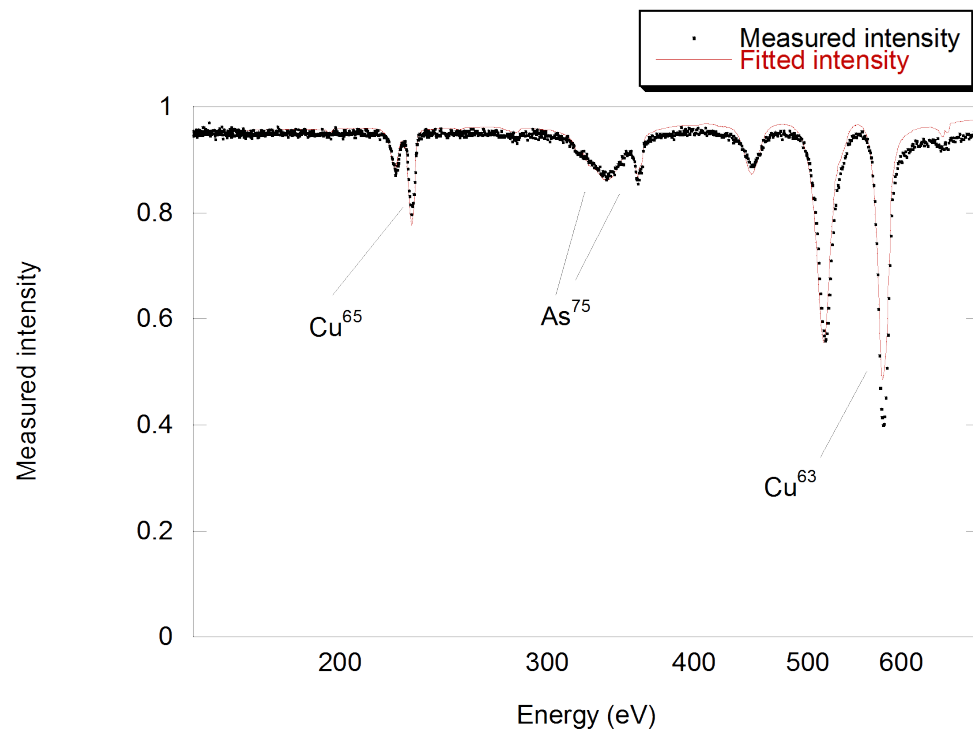
REFIT is a toolkit for analysis of transmission (and capture) neutron resonances spectra.

Calculation of
Doppler broadening
(various models)

energy dependence of
the nuclear cross-
section from the
resonance parameters

Calculation of
resolution


Fitting of the simulated spectra with
experimental data to find $n(x)$



Element	Certified %wt	Estimated %wt
As	0.194	0.20
Sn	7.2	7.65
Zn	6.02	6.97
Sb	0.5	0.45
Mn	0.2	0.21
Ag	-	1.2×10^{-3}

Simplified NRT analysis approach: area calibration

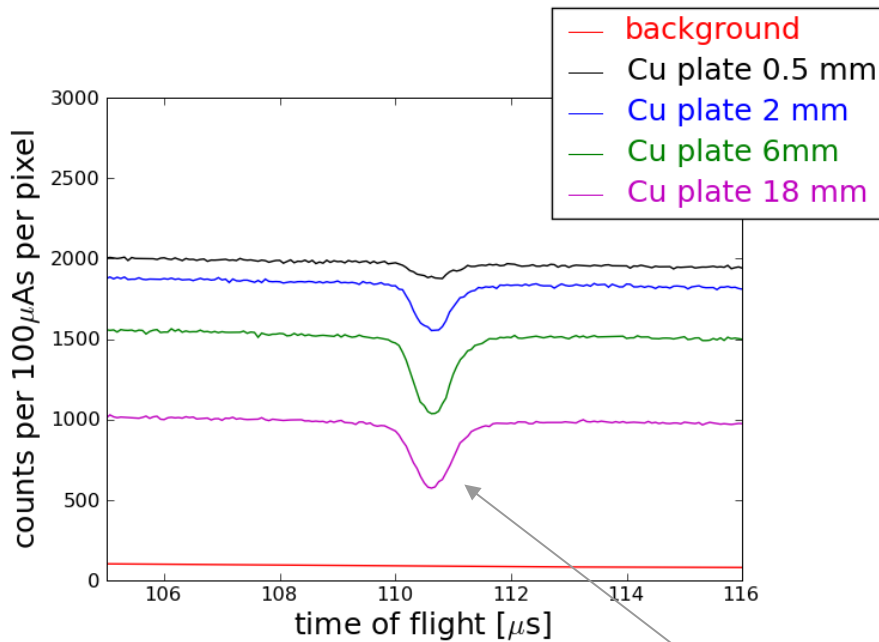
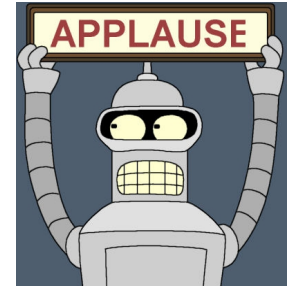
The optimum approach is ideal but quite long (both in terms of machine time and operator time)



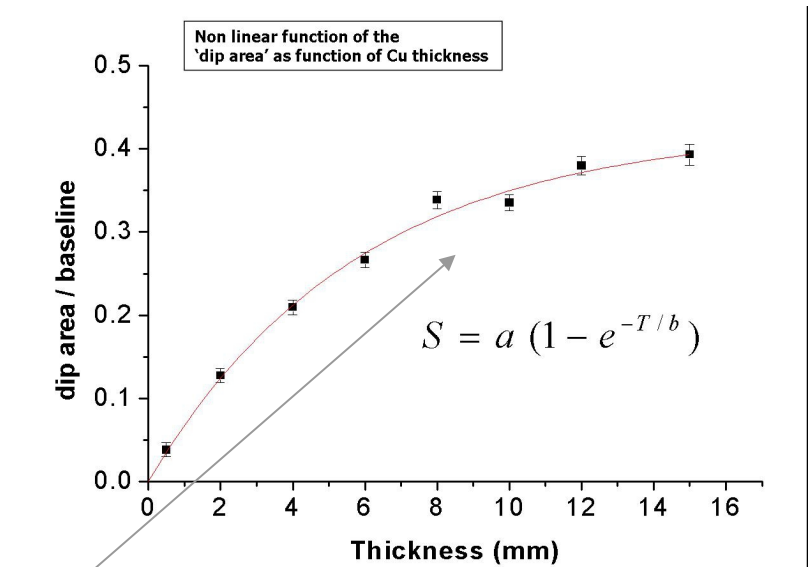
Images require the analysis of hundreds of spectra.....

Simplified NRT analysis approach: area calibration

A suitable series of calibrated samples (for instance metal plates) can lead to an approximated but much simpler analysis



Saturation effect



And what happens once the neutrons are adsorbed?

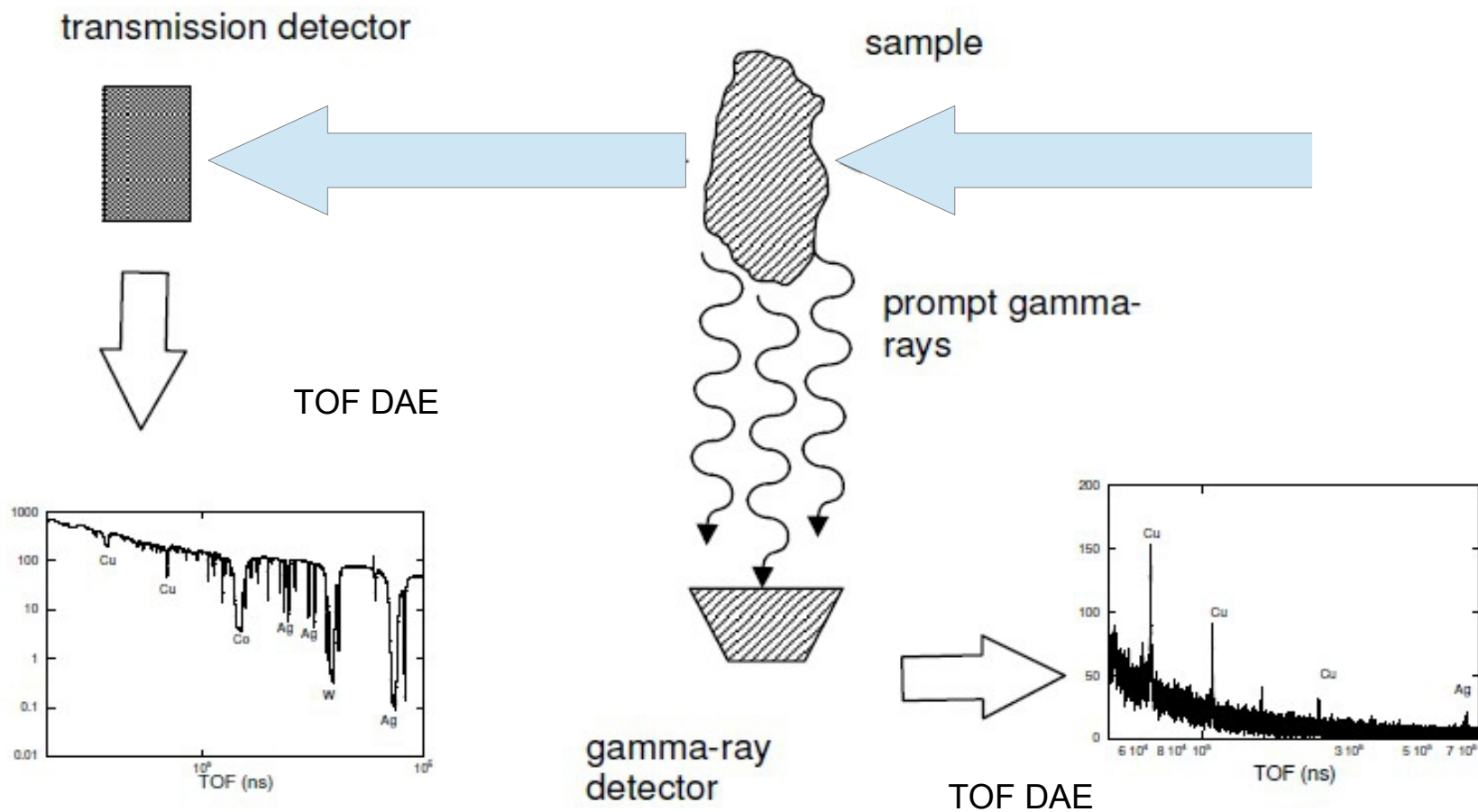


After neutron capture, most (not all....) isotopes deexcite via a PROMPT gamma emission

Prompt means “with characteristic times of the order of ps or less”

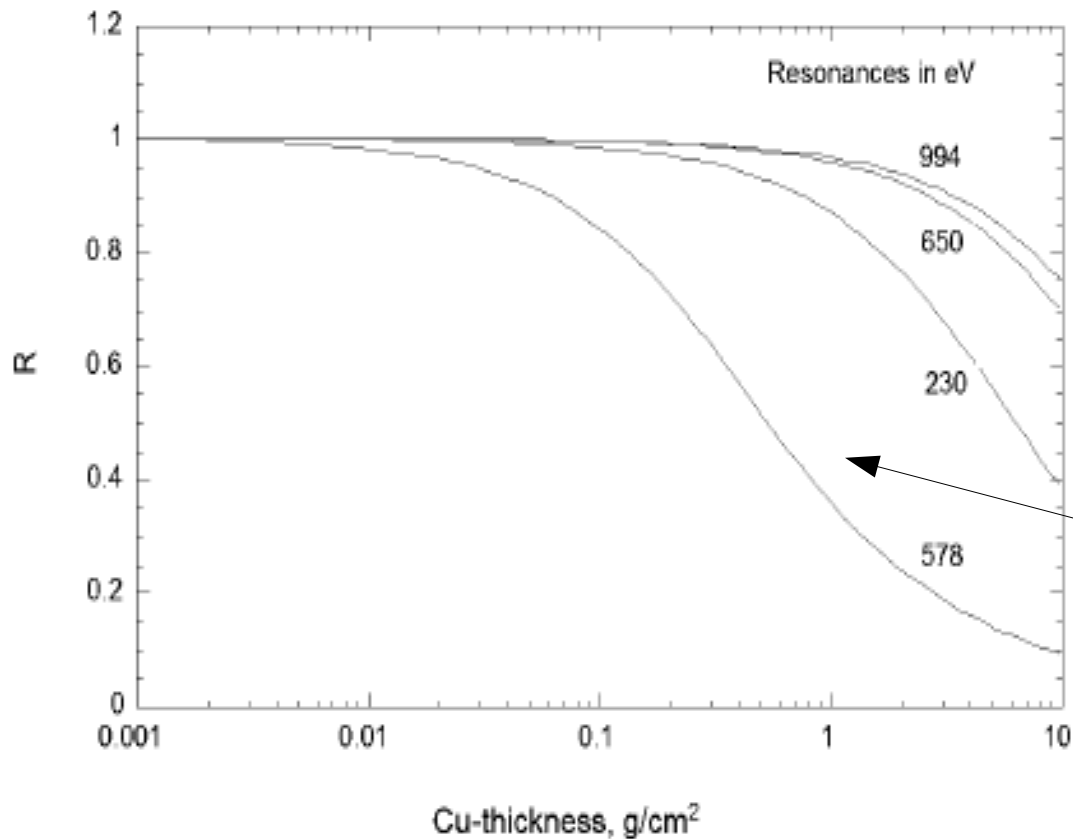
From the point of view of epithermal neutrons, that means IMMEDIATLY

Thus the time information is preserved



Again, a fully quantitative approach is not simple.....

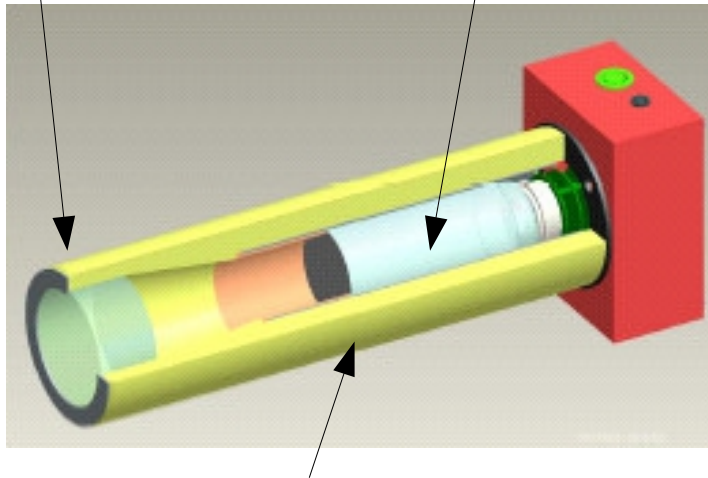
But NRCA has the advantage that a gamma detector is always simpler than a neutron detector!



Advanced set-up for NRCA

YAP crystal

PMT



Lithium carbide housing

YAP is the ideal detector for NRCA

FAST

Neutron-insensitive



YAP scintillator:

51 mm \varnothing , 25 mm thick

Array of 28 γ -detectors

~20% solid angle coverage

Application to Cultural Heritage



Selection of bronze axes from GIA University of Groningen (the Netherlands). H. Postma *et al.* 2005

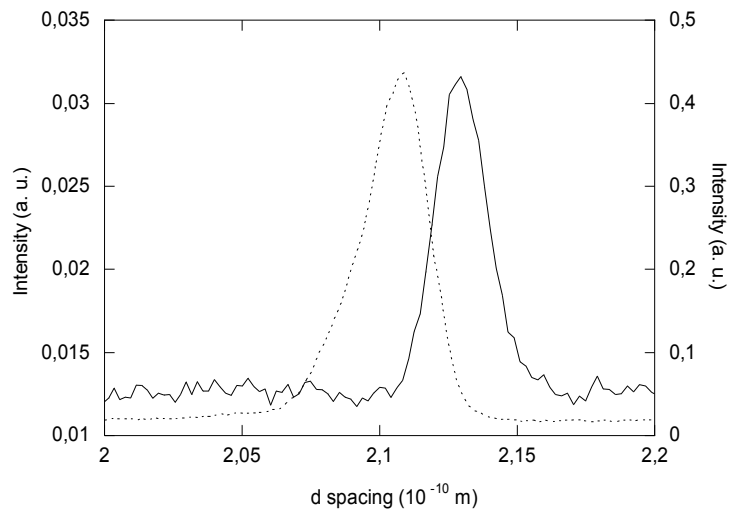
Non-destructive material analysis of findings and museum objects

axe	Cu	Sn	As	Sb	Ag	Fe	Co	Zn	Pb	In (ppm)
1	100*	9.42 ±0.11	0.0774 ±0.0011	0.0736 ±0.0010	0.1004 ±0.0010	≤0.32	nu.l.	0.100 ±0.011	≤1.3	17.4 ±0.3
2	100	15.68 ±0.19	0.292 ±0.004	0.0727 ±0.0010	0.0143 ±0.0005	0.50 ±0.02	0.067 ±0.002	0.224 ±0.014	≤1.2	23.2 ±0.4
3	100	4.96 ±0.06	0.183 ±0.004	0.355 ±0.005	0.153 ±0.002	≤0.03	0.052 ±0.003	≤0.05	≤2.4	7.5 ±0.4
4	100	3.79 ±0.06	0.189 ±0.027	0.0395 0.0008	0.0076 ±0.0007	≤0.03	nu.l.	≤0.09	≤2.3	4.1 ±0.7
5	100	0.082 ±0.015	1.012 ±0.018	2.225 ±0.045	0.662 ±0.005	≤0.08	nu.l.	≤0.29	≤2.2	4.4 ±1.0

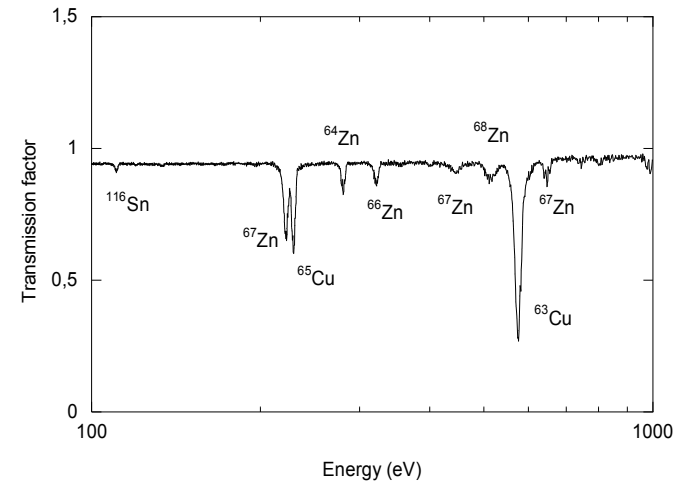
Application to Cultural Heritage

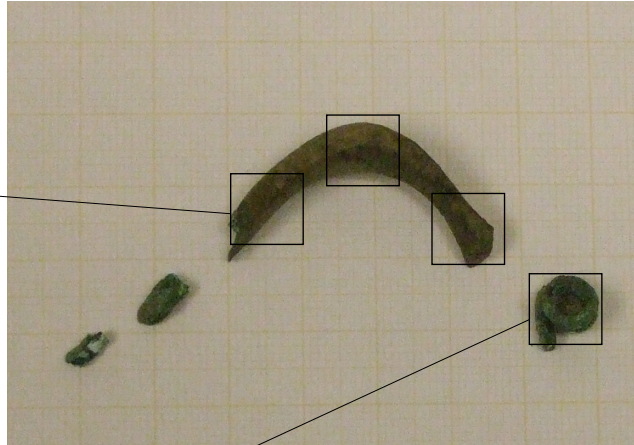
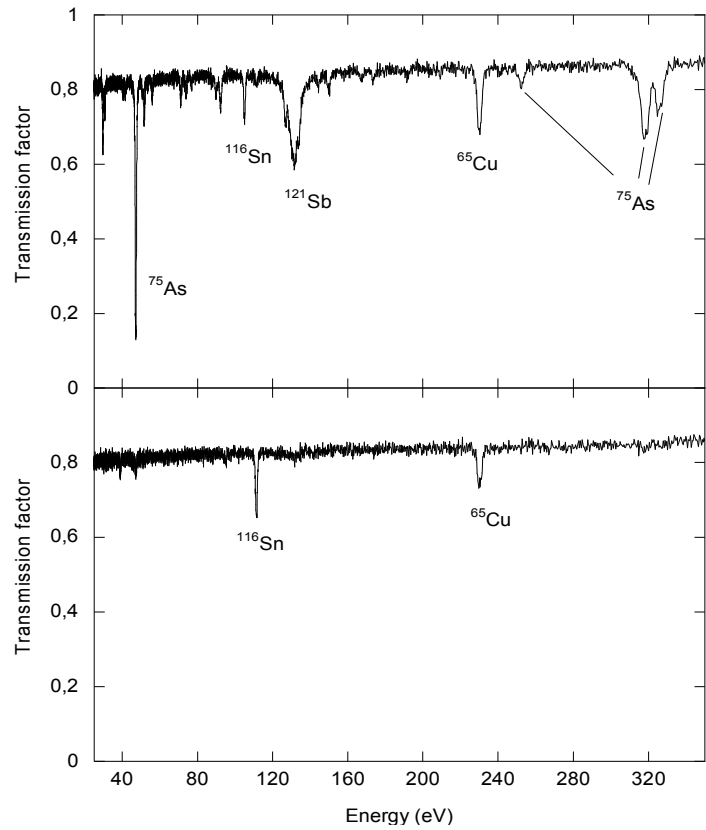
Natural complement to Vegard's law

$$a = a_A^0 (1 - X_B) + a_B^0 (X_B)$$

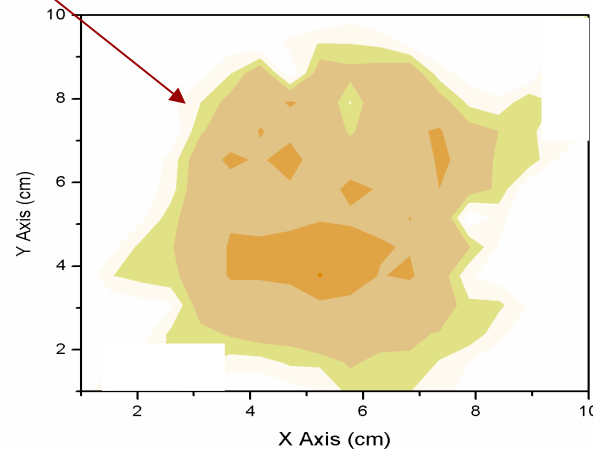
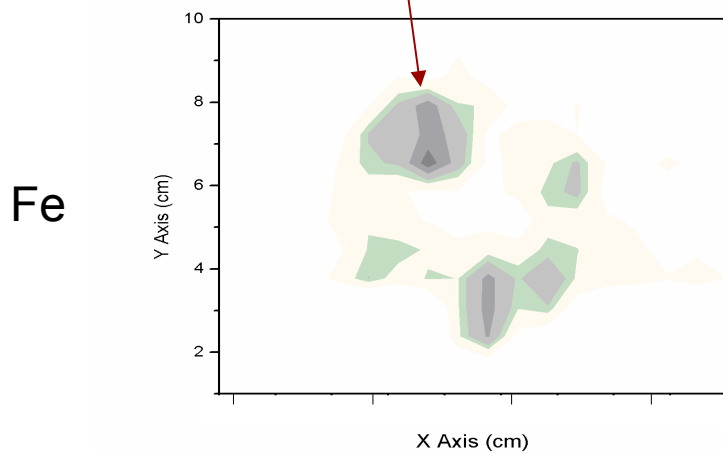
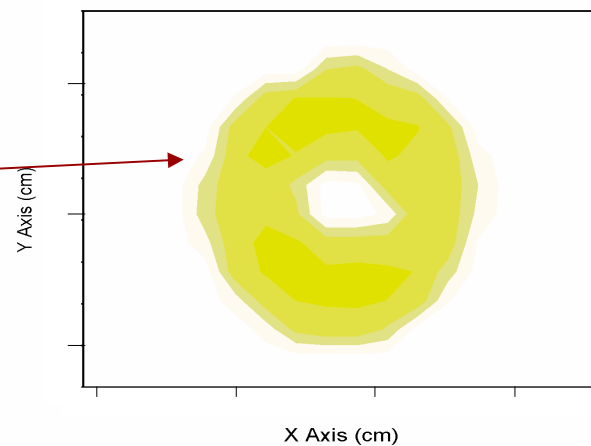
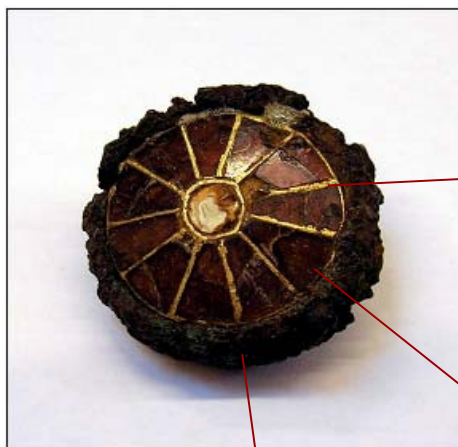


...but this just in the case of binary alloys.....

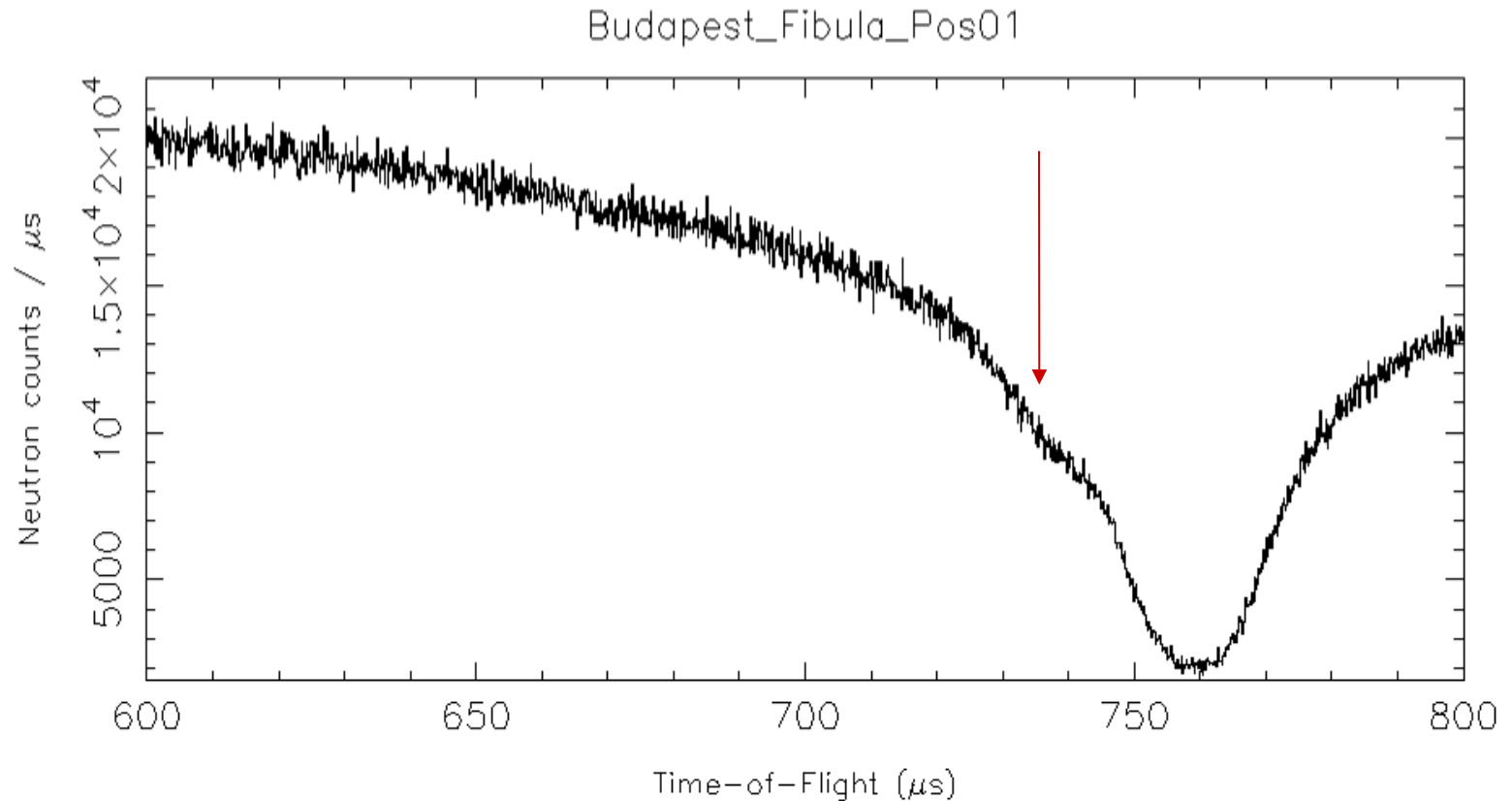




Fibula VI - VII century C. E. From the Kölked-Feketekapu site (National Hungarian Museum, Budapest)



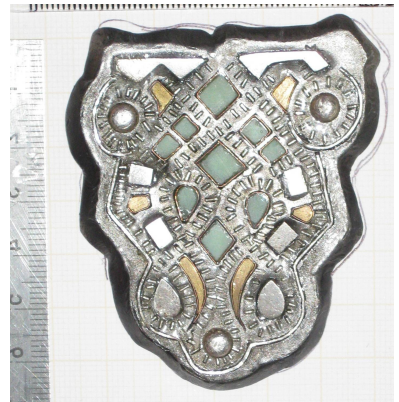
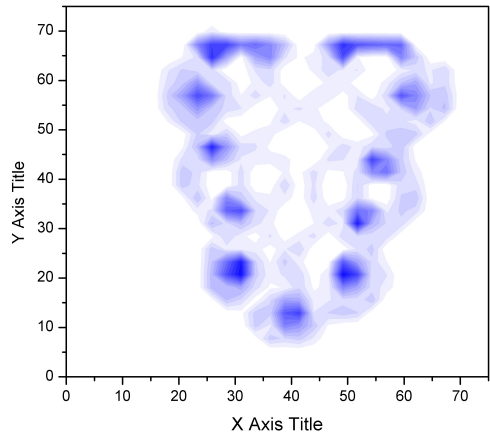
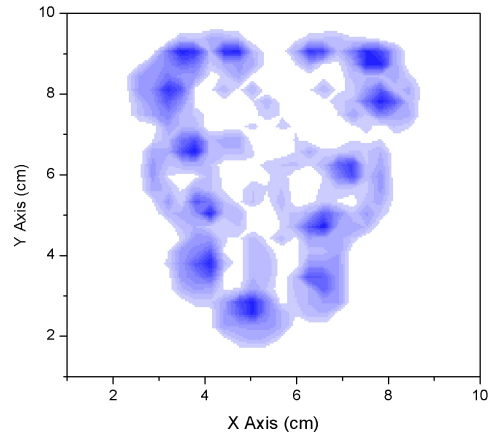
The distribution of gold “follows” the one of silver (the 5.1 eV resonance of silver is always visible as an edge of the 4.9 eV resonance of gold).



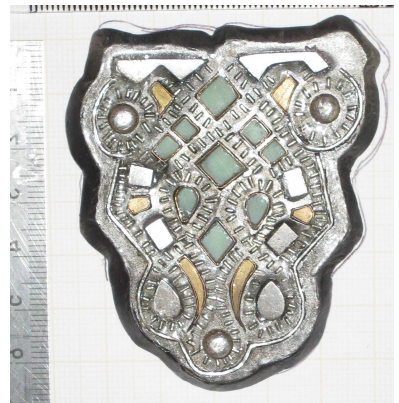
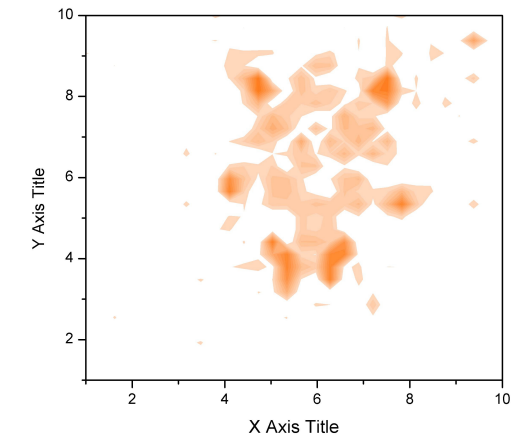
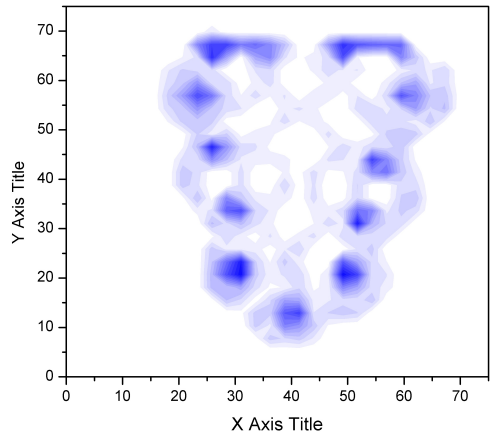
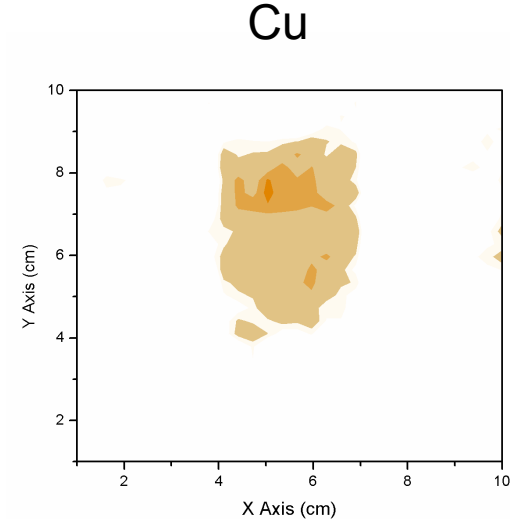
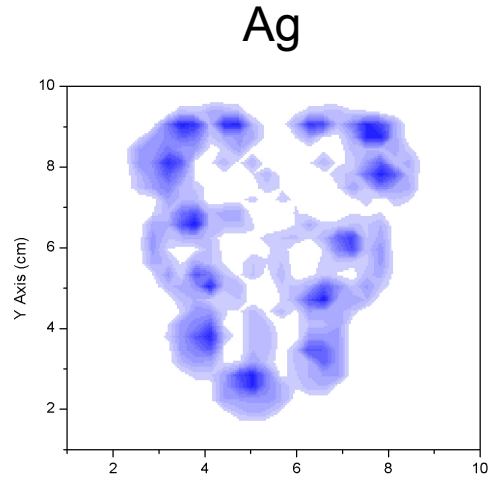
Conclusion: silver is significantly present in the gold alloy of the inlaying but not into the copper alloy of the base plate.

Belt mount VII century C. E. From the Kornye site (National Hungarian Museum, Budapest)

Ag

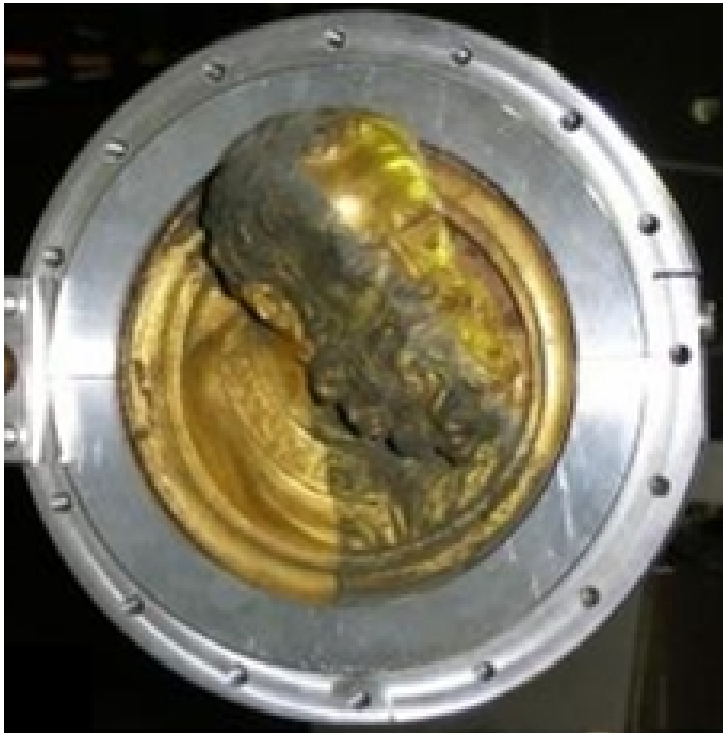


Borchia di cintura VII sec. d. C. dal sito di Kornye (Museo Nazionale Ungherese, Budapest)



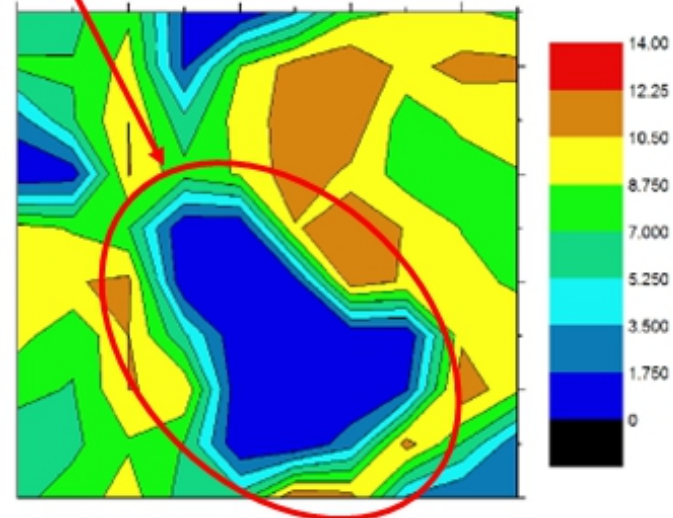
Lorenzo Ghiberti, Heads of Prophets, from the gates of Florence baptistry (Opificio delle Pietre Dure, Firenze)

Gold glided bronze





In the North gate head, the gold leaf was completely removed by the effect of time from the most exposed parts (nose, eyebrows)





On the other hand, in the head from the “Gate of Heavens” the gold leaf is still present in exposed parts (beard) even under a heavy patina.

