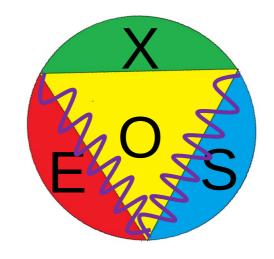




New perspectives for high precision X-ray detection experiments

INFN-CSN5

Young Researcher Grant 2015, n. 17367/2015.



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K- mass precision measurement @ DAФNE

WEIGHTED AVERAGE 493.664±0.011 (Error scaled by 2.5)

The main disagreement is between the two most recent and precise measurements (x-ray energies from kaonic atoms):

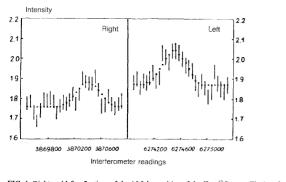


FIG. 1. Right and left reflections of the 4f-3d transition of the K^- - 12 C atom. The interferometer readings are plotted along the abscissa; the detector count rate per 10^{12} protons is plotted along the ordinate. The vertical lines are the experimental values with the corresponding error; the heavy points are the results of a fit.

m_{κ} =493.696±0.007 MeV

A.S. Denisov et al. JEPT Lett. 54 (1991)558

K⁻¹²C, crystal diffraction spectrometer (6.3 eV at 22.1 keV), 4f-3d

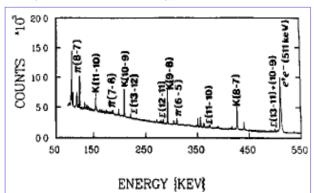
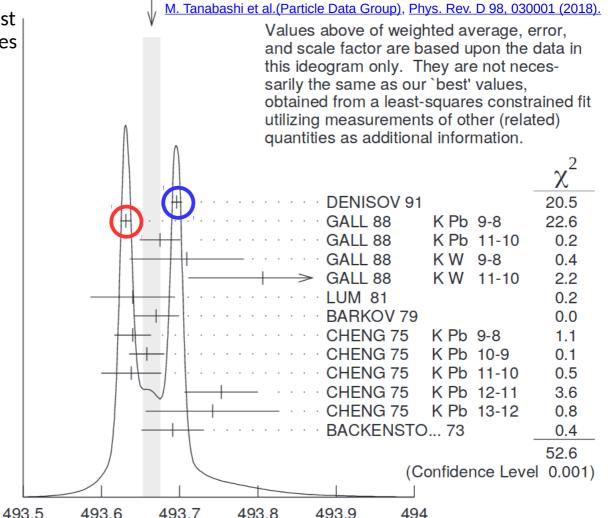


FIG. 1. Untagged Pb x-ray spectrum showing intense kaonic x-ray transitions.



m_K=493.636±0.011 MeV

K.P. Gall et al.

Phys. Rev. Lett. 60 (1988)186

K-Pb, K-W; HPGe detector (1 keV), K-Pb (9 -> 8),

K-Pb (11 -> 10), K-W (9 -> 8), K-W (11 -> 10),



Project's goal



- High resolution (few eV) measurements of the X rays (2-20 keV) emitted in various processes is strongly demanded in: particle and nuclear physics, fundamental science, astrophysics, biology, medical and industrial applications
- Additionally, for some applications (like exotic atoms measurements) X-ray detector systems have to be operated in high background environment.
- These X-rays not always are produced by a point-like source; it is mandatory to develop detectors working with 'extended' (diffused) sources.

VOXES's goal: to develop, test and qualify the first prototype of ultra-high resolution and high efficiency X-ray spectrometer in the range of energies 2 - 20 keV using HAPG bent crystals able to work with 'extended' sources





State of art: high resolution X-ray detectors

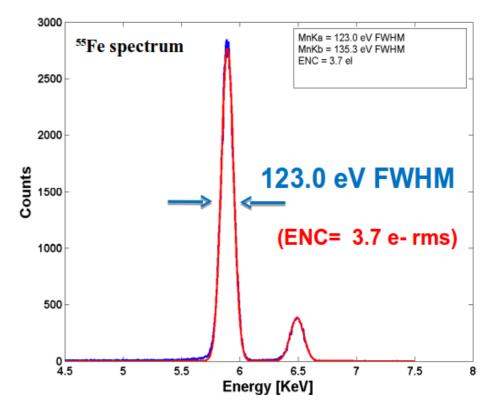


Commonly used detectors for X-rays in the range 1-20 keV are the Solid State Detectors (CCD, SDD, etc...)

However...

The solid state detectors have intrinsic resolution (FWHM ~ 120 eV at 6 keV) given by the electronic noise and the Fano Factor

$$\sigma = \frac{FWHM}{2.35} = \omega \sqrt{W_N^2 + \frac{F \times E}{\omega}}$$



Presently, to achieve ~ eV resolution, two options are available:



- Transition Edge Sensors (TES)
- Crystals and position detectors (Bragg spectrometers)

TES microcalorimeter

a thermal detector measuring the energy of an incident Transition Edge Sensor x-ray photon as a temperature rise (= E/C ~ 1 mK) $R_0\sim 50 \text{ m}\Omega$ Thermometer Decay time constant X-ray energy: $= C / G (\sim 500 \mu s)$ 8.0 Absorber Heat capacity: C Resistance particle Thermal conductance: G absorption Low temperature heat sink Rapid increase in resistance TES microcalorimeter bias point 0.2 Tiny temperature rise $T_{c} \sim 50 \text{ mK } !!!$ 119 120 121 Temperature (mK) pixel ~ 350 x 350 µm²

LIMITATIONS:

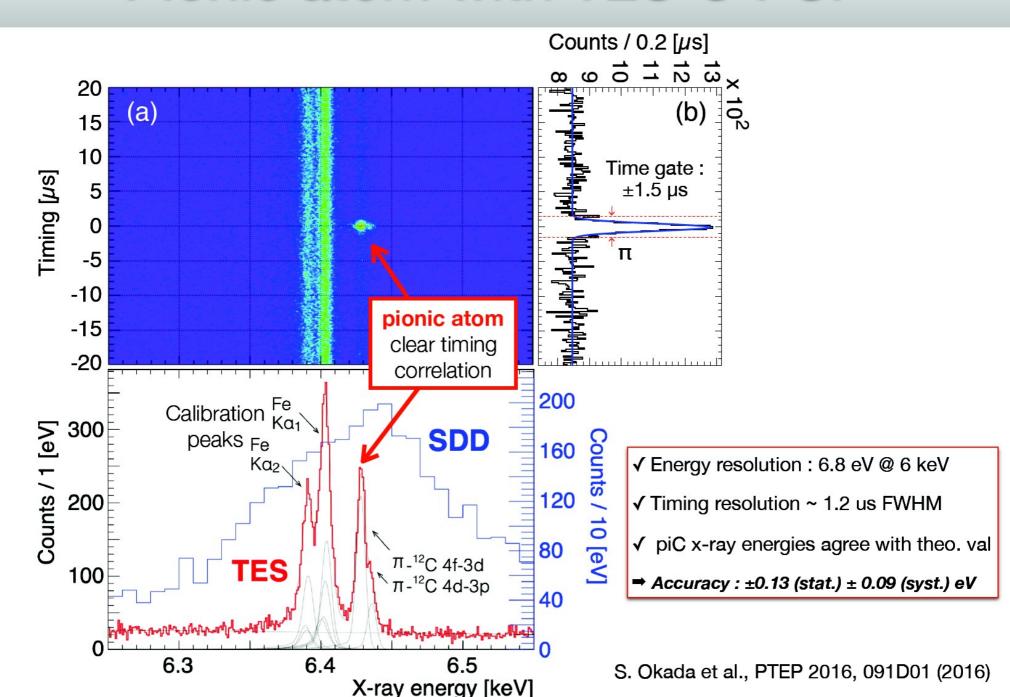
- not optimised for E < 5 keV
- very small active area
- prohibitively high costs
- complex cryogenic system needed
- complex calibration

high energy resolution ($\Delta E / E \sim 10^{-3}$)

TES: ΔE (FWHM) ~ 5 eV @ 6 keV X-ray (ref. SDD: ΔE (FWHM) ~ 150 eV @ 6 keV)

Reference: Bennet et al., Rev. Sci. Instrum. 83, 093113 (2012)

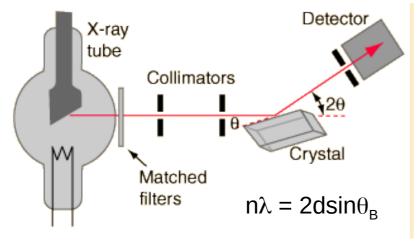
Pionic atom with TES @ PSI





Crystal spectrometers limitations





High resolution can be achieved depending on the quality of the crystal and the dimensions of the detectors

Geometry of the detector determines also the energy range of the spectrometer

But....

Crystals response may not be uniform (shape, impurities, ecc.)

Lineshapes are difficult to be measured within few eV precision (surface scan)

In accelerator environments particles may hit the detector

Background reduction capability is mandatory

Typical d (Si) $\approx 5.5 \text{ Å}$

 $\theta_{\rm B}$ < 10° for E > 6 keV (forward & difficult)



Limitation in efficiency



Mosaic crystals

Mosaic crystal consist in a large number of nearly perfect small crystallites.

Mosaicity makes it possible that even for a fixed incidence angle on the crystal surface, an energetic distribution of photons can be reflected

Increase of efficiency (focusing) ~ 50

Loss in resolution

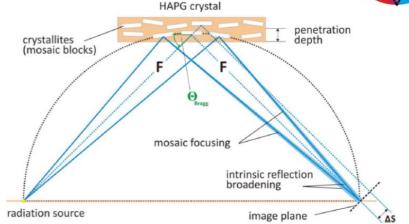
(focusing) ~ 50

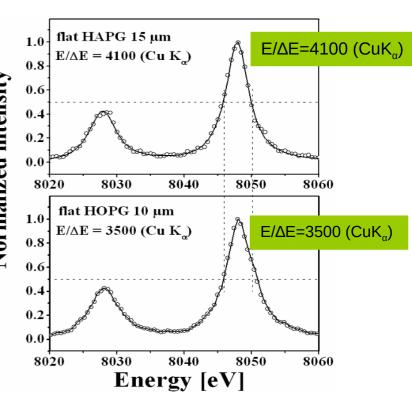
Pyrolitic Graphite mosaic crystals (d = 3.354 Å):

Highly Oriented Pyrolitic Graphite (HOPG, $\Delta\theta \approx 1^{\circ}$)

Highly Annealed Pyrolitic Graphite (HAPG, $\Delta\theta \approx 0.07^{\circ}$)

flexible HAPG has twice higher spectral resolution, while flexible HOPG approximately twice higher reflectivity





H. Legall, H. Stiel, I. Grigorieva, A. Antonov et al., FEL Proc. 2006

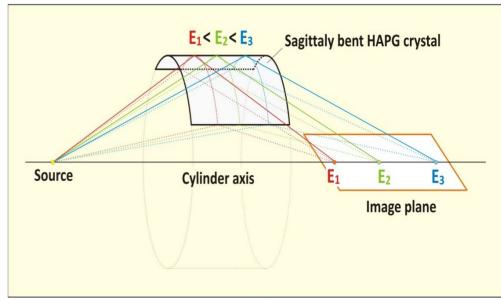




HAPG: High Annealed Pyrolitic Graphite



- Bending does not influence resolution and intensity
- Mosaic spread down to 0.05 degree
- Integral reflectivity $\sim 10^2$ higher than for other crystals
- Variable thickness (efficiency)
- Excellent thermal and radiation stability

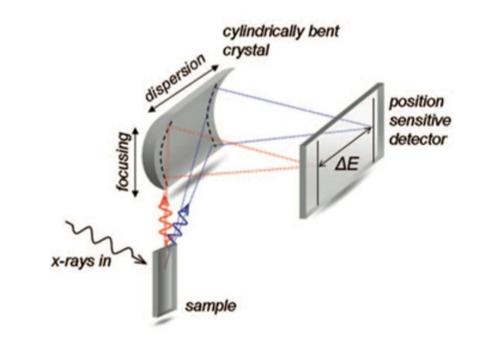


Characterization of HAPG mosaic crystals using synchrotron radiation

Martin Gerlach, ^a Lars Anklamm, ^b Alexander Antonov, ^c Inna Grigorieva, ^c Ina Holfelder, ^a Birgit Kanngießer, ^b Herbert Legall, ^c Wolfgang Malzer, ^b Christopher Schlesiger ^b and Burkhard Beckhoff ^a*

J. Appl. Cryst. (2015). 48

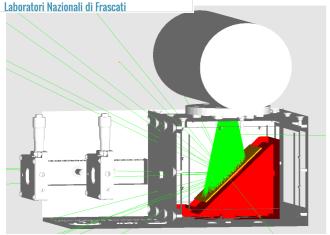


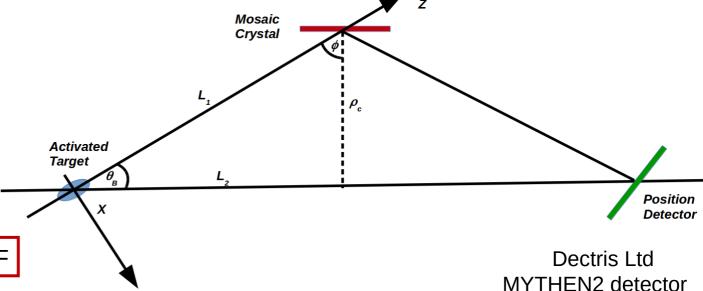




Von Hamos configuration







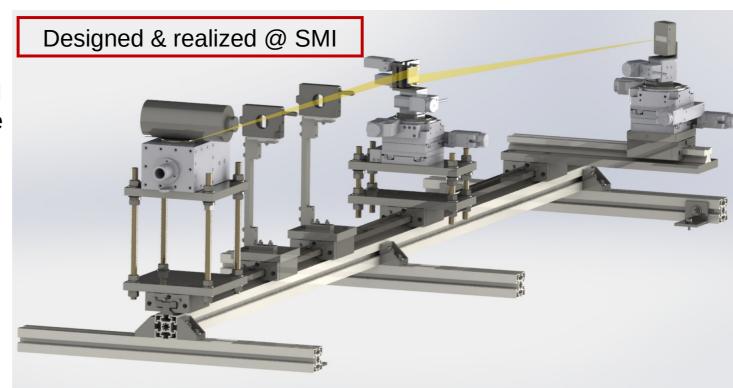
Designed & 3D-printed @ LNF

For a given X-ray energy the Bragg angle ($\theta_{\rm B}$) and the curvature radius of the crystal ($\rho_{\rm c}$) completely determine the position of the source, the crystal and the position detector

$$L_1 = \frac{\rho_c}{\sin \theta_B}$$









The source size problem What has been done so far?



Bragg spectroscopy is usually exploited by XAS, XES, synchrotron light users, etc...

The x-ray source, which was used for the measurements, is a low power microfocus x-ray tube (IfG) with a source diameter of about $50 \ \mu m$ Measurements were performed with the Cu K_{\alpha} emission of a Cu anode at 8 keV. The spec-

III. SPECTROMETER SETUP

The spectrometer consists of three principal components: the X-ray source, the HAPG optic, and the position sensitive detector. As source a watercooled 100 W micro focus X-ray tube with a tungsten anode and a focus size of 50 μ m is used. The emitted radiation is focused onto the sample by a polycapillary full lense with a spot size of 35 μ m. The HAPG

This means:

- Point-like sources
- High yields (no need to increase them)
- 1 eV (Si, Mica, etc) or 2-3 eV (HAPG) resolution is very easily achievable

Laser-produced plasmas were created using the "Phoenix" Nd glass laser (the Lebedev Physical Institute) operated at a wavelength of 0.53 μm with pulse energy up to 10 J and 2 ns pulse duration. The laser beam was focused onto massive Mg, Al, Ti, or Fe targets (see Fig. 2). The focal spot diameter was about $\sim 15~\mu m$

But what if we really need wider sources for higher statistics? Or what if our photons come from a diffused isotropic source?





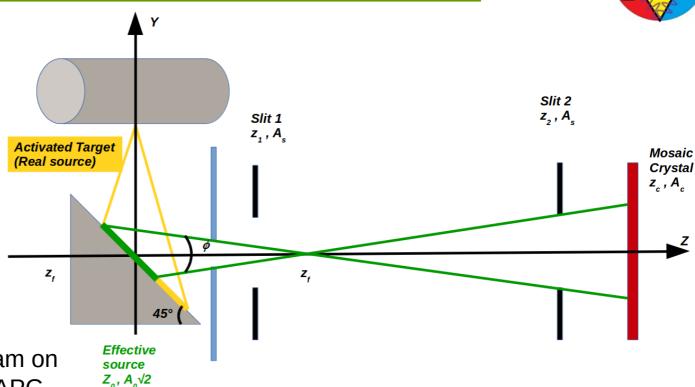
Vertical spread

Setup box front panel

 $z_h = 52,75 \text{ mm}$ $A_h = 5,9 \text{ mm}$



The vertical spread of the X-ray beam is fixed by the slits positions (z_1,z_2) and their frame size (A_s) , together with the hole in the front panel of the setup box (z_h,A_h) .



The vertical spread of the beam on the target (A_0) and on the HAPG crystal (A_c) are then:

$$A_0 = 2z_f t g \phi$$

We first define the position of the intersection point z_f and the ϕ angle :



$$A_c = 2(z_c - z_f)tg\phi$$

$$tg\phi = \frac{A_s + A_h}{2(z_2 - z_h)} \qquad z_f = z_2 - \frac{A_s}{2tg\phi}$$



Characterization @ 6-8 keV

 $S_0' = 0.8 \text{ mm}$

 $S_0 = 0.43 \text{ mm}$

 ρ = 206,7 mm

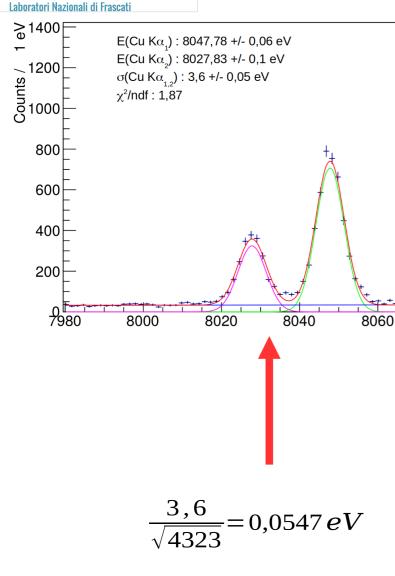
8080 810 X-ray energy (eV)

8100

 $\Delta\theta = 0.57$ °

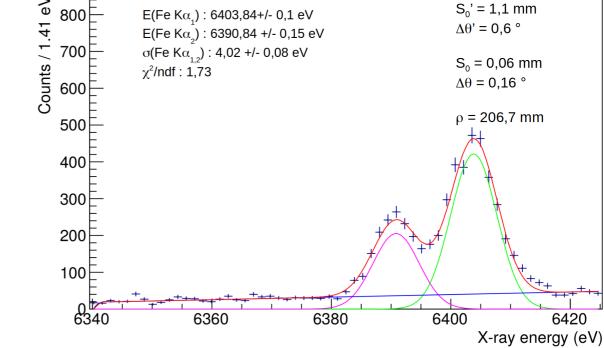
 $\Delta\theta$ ' = 0,6 °





In the limit of a background free pure gaussian peak, the precision is related to the resolution via:

$$\delta E = \frac{\sigma E}{\sqrt{N}}$$





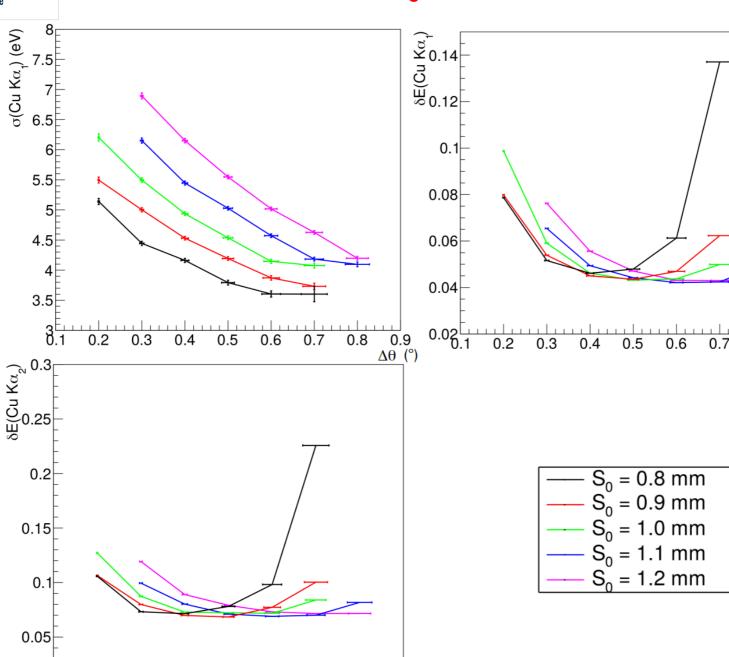


Scan over $\Delta\theta$ ', S_0 ' (Cu 206,7 mm ρ)



0.9

 $\Delta\theta$ (°)



Δθ (°)



0.2

0.3

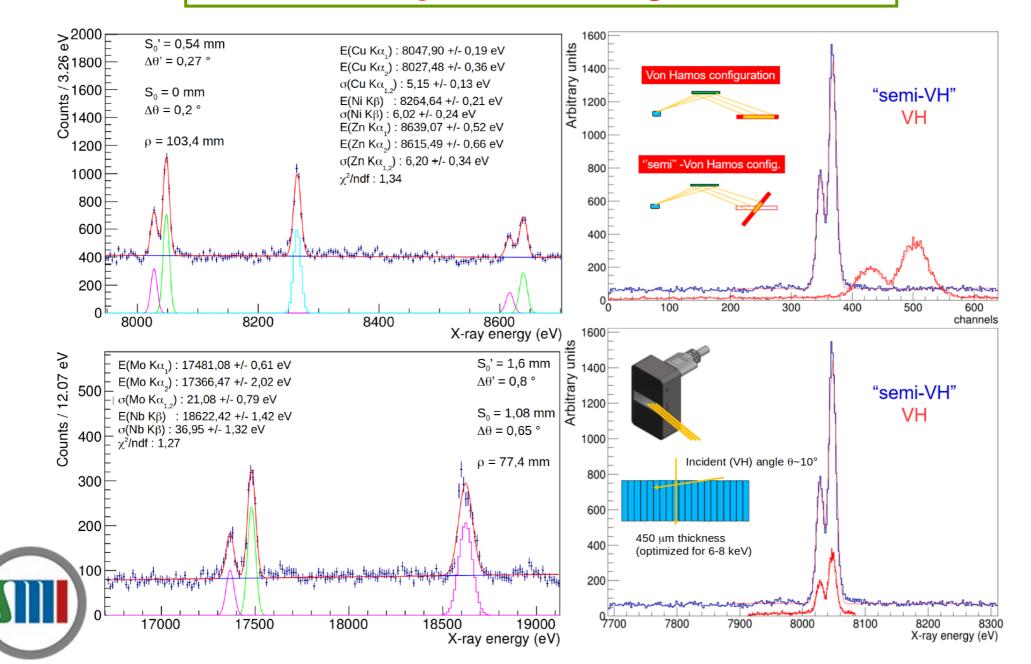
0.5

0.6



Semi VH configuration: wider dynamic range



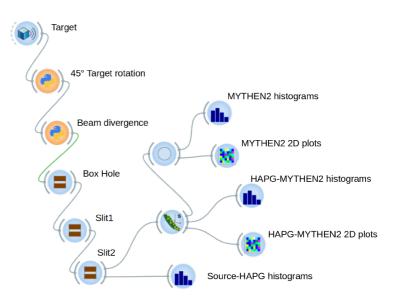




Results summary



Element	$\rho_{c}\left(mm\right)$	Parameter	value (eV)	$S_0'/\Delta\theta' (mm,^\circ)$
	77,5	$\sigma(K\alpha_{1,2}) \ \delta(K\alpha_1) \ \delta(K\alpha_2)$	$4,17 \pm 0,16$ 0,11 0,18	0,3/0,24 0,6/0,44 0,6/0,44
Fe	103,4	$\sigma(K\alpha_{1,2}) \ \delta(K\alpha_1) \ \delta(K\alpha_2)$	$4,05 \pm 0,13$ 0,09 0,13	0,3/0,18 0,7/0,34 0,7/0,34
	206,7	$\sigma(K\alpha_{1,2}) \ \delta(K\alpha_1) \ \delta(K\alpha_2)$	$4,02 \pm 0,08$ 0,1 0,15	$ \begin{array}{c} 1,1/0,60 \\ 1,2/0,70 \\ 1,2/0,70 \end{array} $
	77,5	$\sigma(K\alpha_{1,2}) \ \delta(K\alpha_1) \ \delta(K\alpha_2)$	$6,8 \pm 0,07$ 0,07 0,1	0,3/0,16 0,6/0,32 0,6/0,32
Cu	103,4	$ \frac{\sigma(K\alpha_{1,2})}{\delta(K\alpha_{1})} $ $ \delta(K\alpha_{2}) $	$4,77 \pm 0,05$ 0,04 0,07	0,3/0,16 0,7/0,32 0,7/0,32
	206,7	$\sigma(K\alpha_{1,2}) \ \delta(K\alpha_1) \ \delta(K\alpha_2)$	$3,60 \pm 0,05$ 0,04 0,07	0.8/0.60 $1.1/0.70$ $1.1/0.70$
Cu	103,4	$\sigma(K\alpha_{1,2}) \ \delta(K\alpha_1) \ \delta(K\alpha_2)$	$5,15\pm0,13$ 0,10 0,21	0,5/0,27 0,6/0,22 0,6/0,22
Ni	103,4	$\sigma(Keta) \ \delta(Keta)$	$6,02 \pm 0,24$ 0,13	0,5/0,27 0,6/0,22
Zn	103,4	$\sigma(K\alpha_{1,2}) \ \delta(K\alpha_1) \ \delta(K\alpha_2)$	$6,20 \pm 0,34$ 0,26 0,42	0,5/0,27 0,6/0,22 0,6/0,22
Мо	77,5	$\sigma(K\alpha_{1,2}) \ \delta(K\alpha_1) \ \delta(K\alpha_2)$	$21,1\pm0,8$ $0,6$ $2,0$	1,6/0,80 $1,6/0,80$ $1,6/0,80$
Nb	77,5	$\sigma(K\beta) \ \delta(K\beta)$	$36,9 \pm 1,3$ $1,3$	1,6/0,80 1,6/0,80



To be checked with X-ray tracing simulations

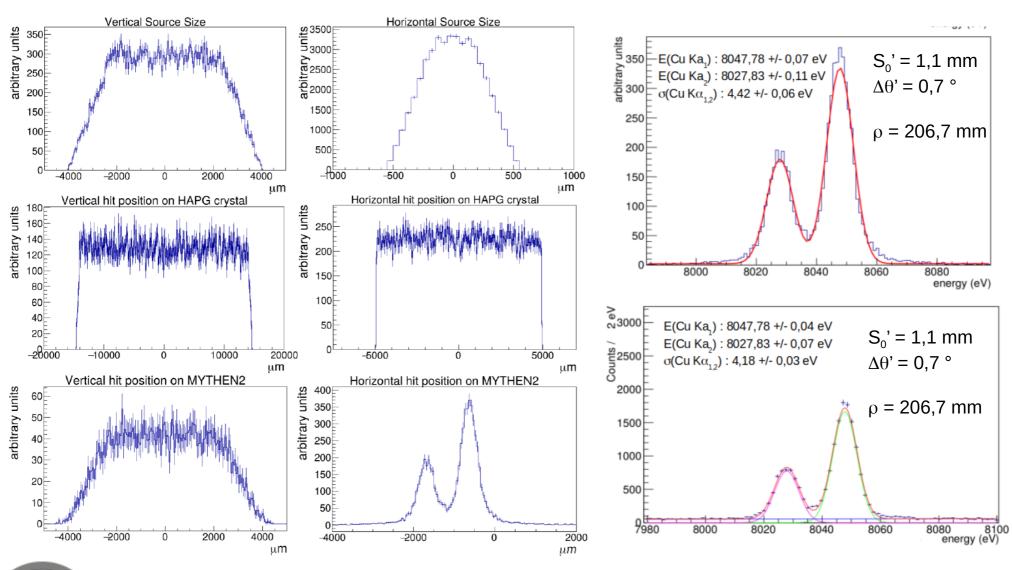
Shadow and XOP software integrated in the OASYS environment.





Ray tracing simulations









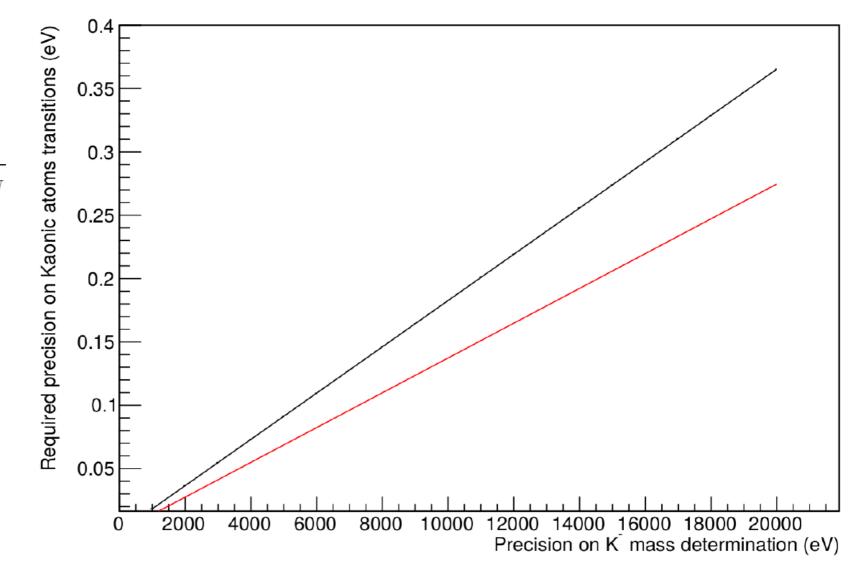
K mass determination



$$E_{X\to Y}^K = \frac{Z^2 e^4 \mu_{KN}}{8h^2 \varepsilon_0^2} \left(\frac{1}{Y^2} - \frac{1}{X^2}\right) \qquad \qquad \sigma m_K = \frac{m_K^2}{\mu_{KN}^2} \frac{1}{Z^2} \frac{10^6}{26, 6} \frac{\sigma E_{X\to Y}^K}{\left(\frac{1}{Y^2} - \frac{1}{X^2}\right)}$$

$$\mu_{KN} = \frac{m_K m_N}{m_k + m_N}$$

$$m_K = \frac{\mu_{KNm_N}}{m_N - \mu_{KN}}$$





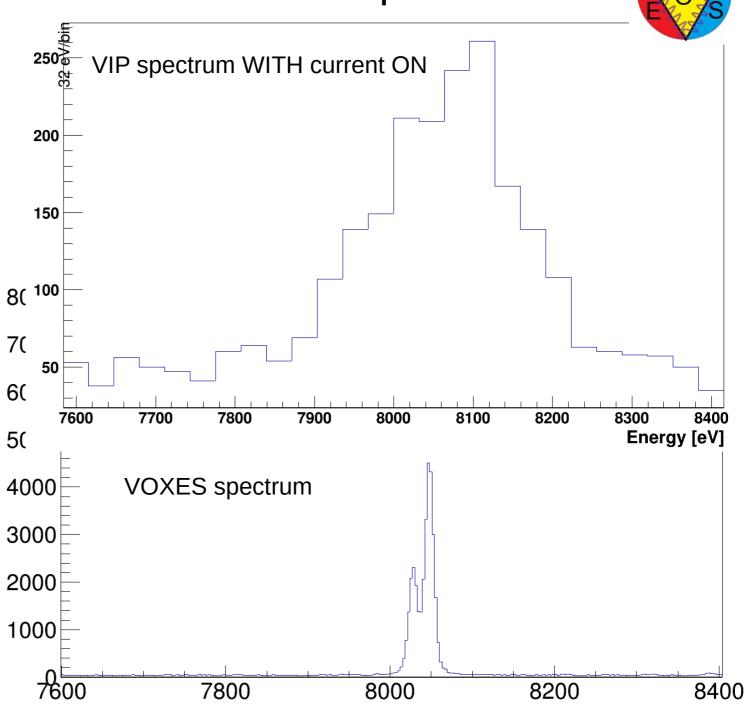


What about the VIP Cu spectrum?

How much could one gain with a \approx 100 times better resolution?

Estimations including efficiency and MC to optimize a possible setup are needed

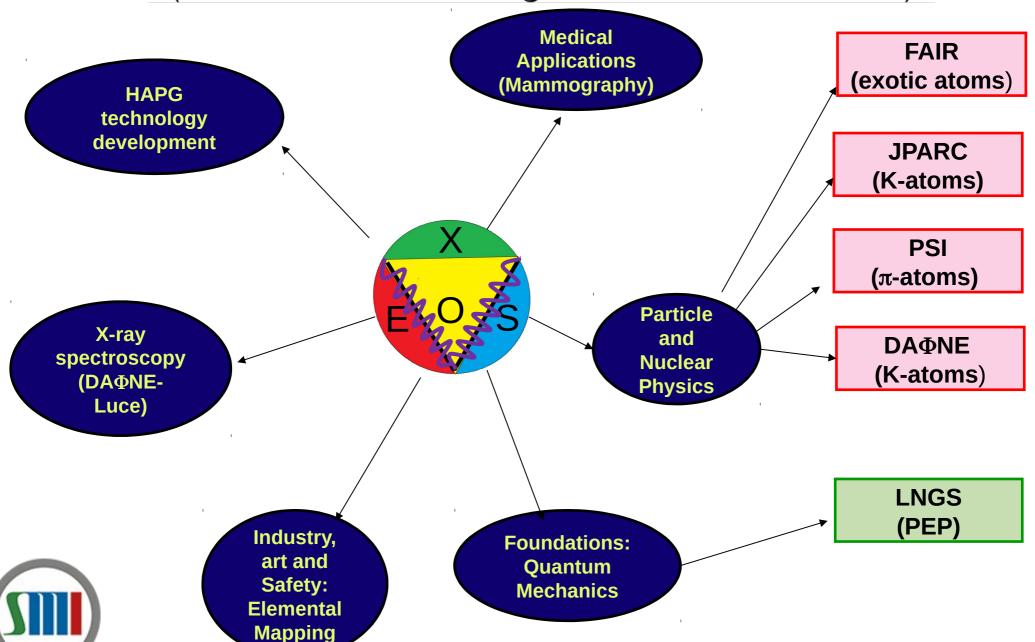






Impact (scientific, technological, socioeconomic)







Impact (scientific, technological, socioeconomic)



Transportable and Agile Spectrometer for metal Trace in Edible liquids : TASTE

INFN-CNTT commision "Research 4 Innovation 2020" awarded

Istituto Nazionale di Fisica Nucleare Trasferimento Tecnologico

Hints for possible measurements???



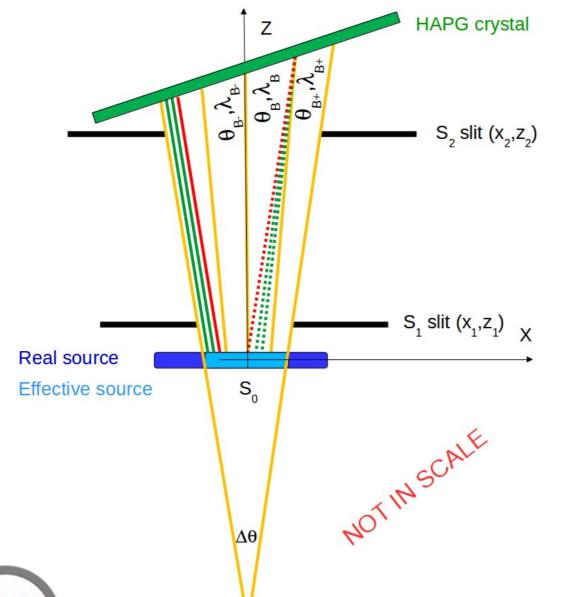


Thanks for your attention



The source size problem





Virtual point-like source

X ray source
Slits
HAPG
Signal photons
Background (?)

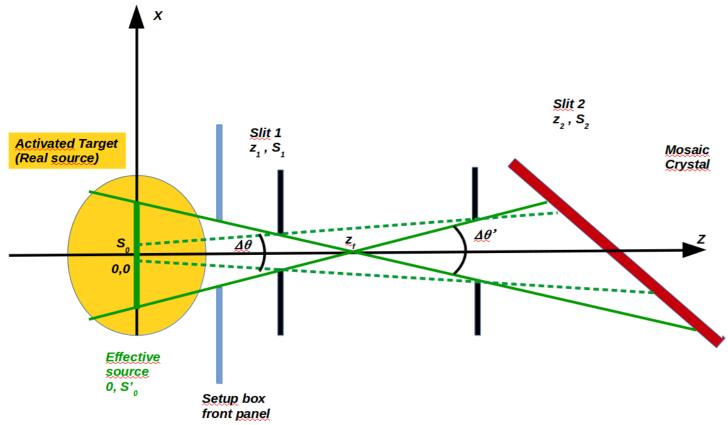
Is it all? Are the red lines really (only) background?

With this configuration you are limited to few tens of microns sources and very compact spectrometers....



Horizontal spread





For each $\Delta\theta$ ', S_0 ' pair, the 2 values of the slits can be found; first, we define the position of the intersection point z_f :

$$z_f = \frac{S_0'}{2} ctg\left(\frac{\Delta\theta'}{2}\right)$$

Then, the 2 slits aperture are defined by:

$$S_1 = \frac{z_f - z_1}{z_f} S_0'$$

$$S_2 = \frac{z_2 - z_f}{z_f} S_0'$$



$$S_c = \frac{z_c - z_f}{z_f} S_0'$$

 $z_{h} = 52,75 \text{ mm}$ $S_{h} = 5,9 \text{ mm}$

