

Young Scientist Workshop
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Germanium Detectors and Natural Radioactivity in food

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GeDet group



→ Introduction to Germanium detectors

- radiation detection in semiconductors
- properties of Germanium detectors

→ Experimental setup

- background measurement
- weak sources
- background reduction

→ Data analysis

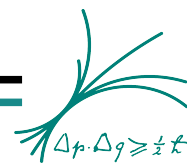
- calibration with KCl salt
- measurement of food rich in Potassium

→ Conclusions

- summary
- outlook



Introduction to Germanium detectors



- RADIATION

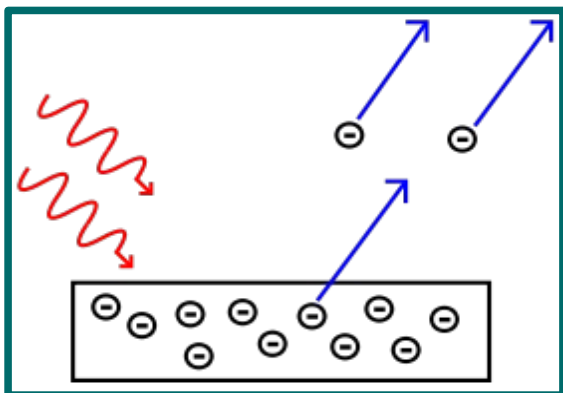


a) which kind of radiation?

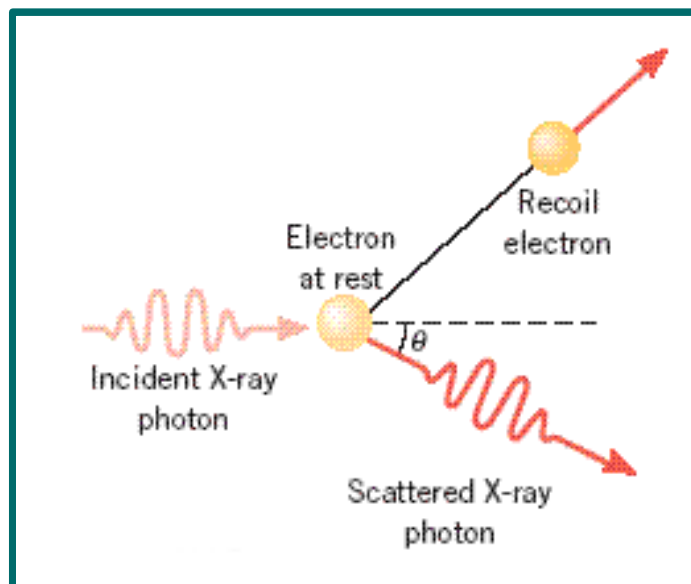
PHOTONS: produced in the decays of radioactive isotopes

b) what can they do in matter?

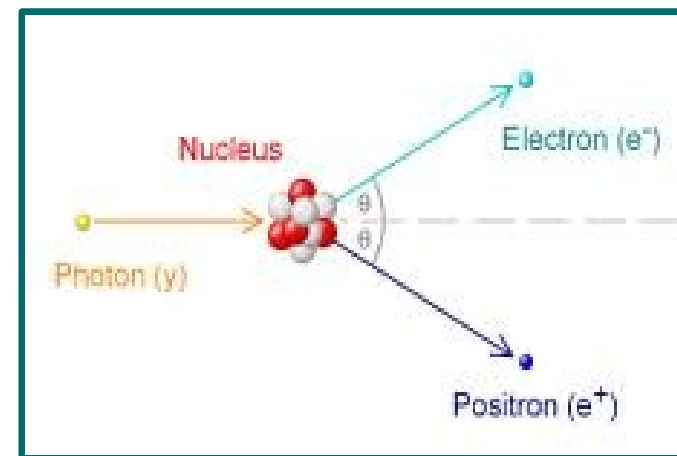
PHOTOELECTRIC EFFECT



COMPTON EFFECT

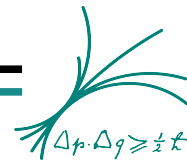


PAIR PRODUCTION



We will use TOTAL ABSORPTION EVENTS

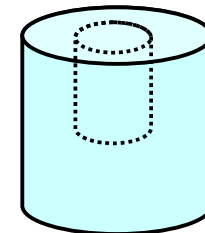




- DETECTOR

a) which kind of detector?

**Closed-ended Coaxial
eXtended Range Germanium detector**

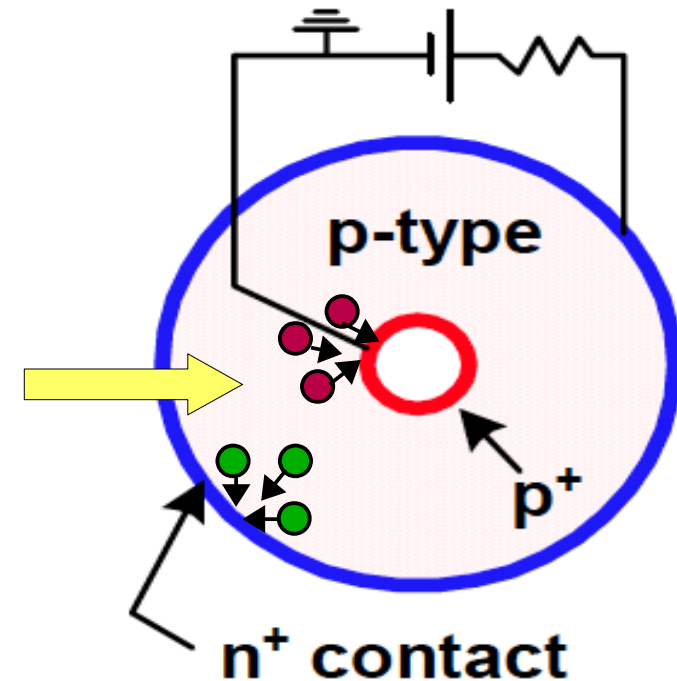


p-type
crystal

b) how does it work?

- **large diode** with a **reverse bias**
→ a **POSITIVE** outside potential

- **radiation** goes into the crystal
→ electron-hole pairs
→ under Electric Field
 - ● electrons go to **n+** contact
 - ● holes go to **p+** contact

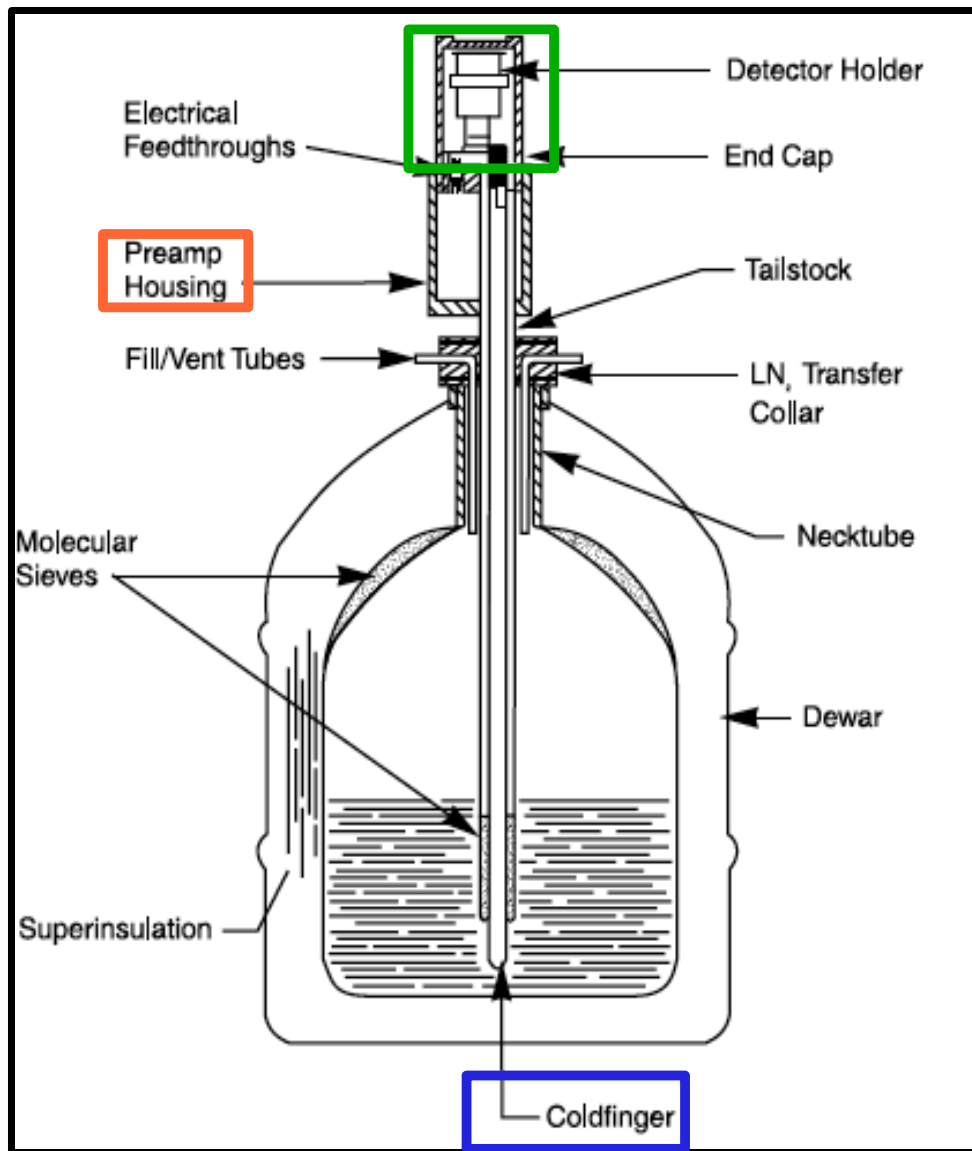


How can we use such a detector?



Introduction to Germanium detectors

$$\Delta f \cdot \Delta g \geq \frac{1}{2} k$$



- A) The detector is in a **clean vacuum chamber**
- B) The detector **has to be cooled**:
- reduce thermal charge carrier generation (**noise**)
 - thermal contact with **LN₂**
- C) charges collected on the **electrodes**
- charge sensitive **preamplifier**
 - sample resulting **pulse**

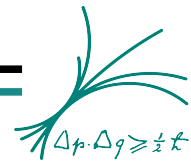


Pulse amplitude proportional to energy deposited in the detector!!

What can be measured? ENERGY SPECTRA

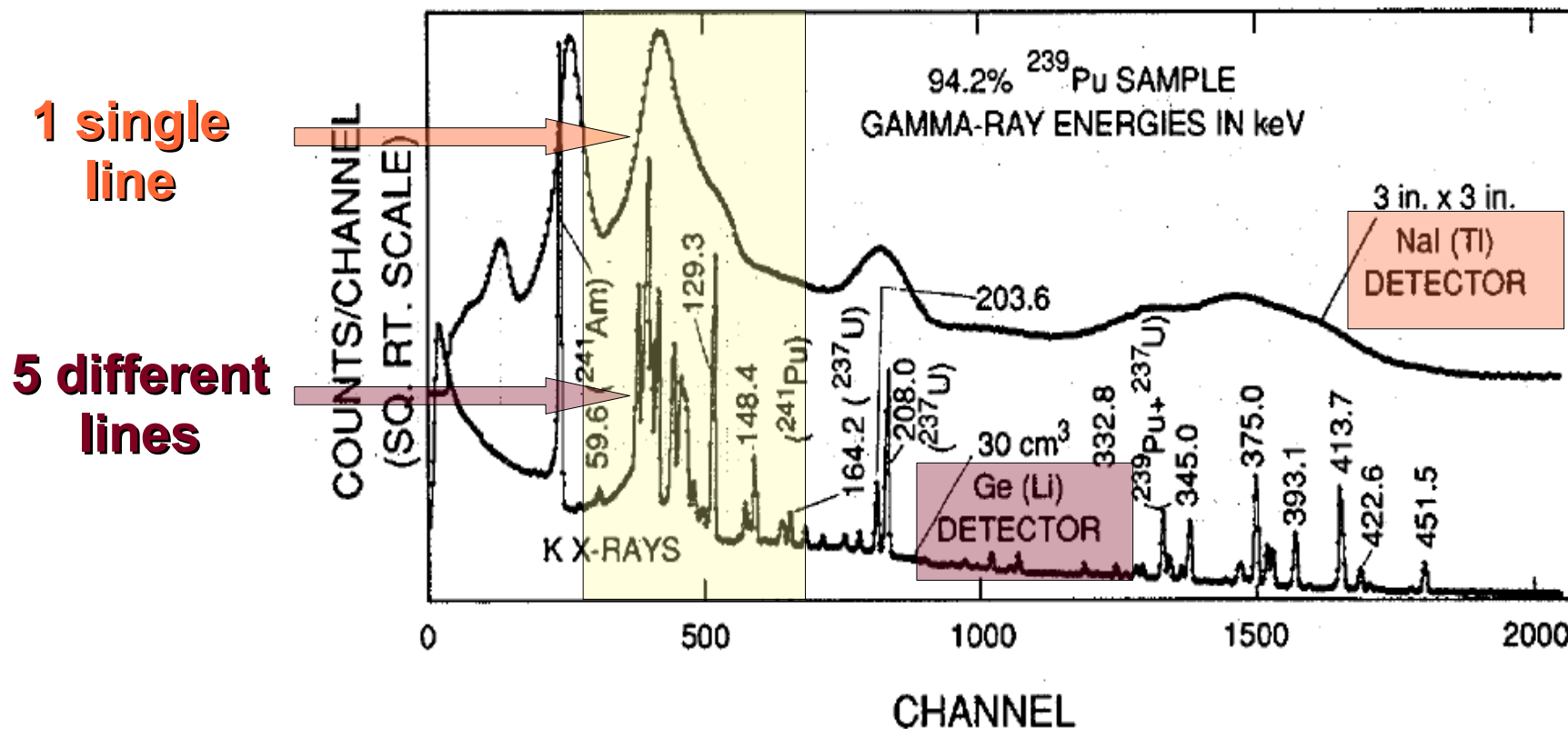


Introduction to Germanium detectors



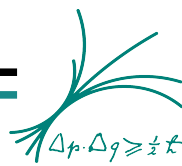
Features of Germanium detector → effect on the spectrum

- small energy required to create e-h pair → large output signal
- ultra pure material → intrinsic detection efficiency close to 100%
- best possible energy resolution



- Resolution is better than expected from statistics (revealed on the full energy peak)





OBSERVED VARIANCE = F * POISSON VARIANCE

→ the total number of **IONIZATION** has a constraint
THEY are **not INDEPENDENT** anymore

NO POISSONIAN STATISTICS

First evaluation of the variance

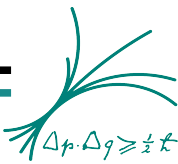
U. Fano, Phys. Rev. 72 (1947) 26

- **energy and momentum conservation**
- the energy deposited is a **FIXED** value E_0
 - fluctuating **BUT** not independent variables
 - N_x lattice excitation
 - N_Q charge carriers

$$\begin{aligned} \epsilon_i &= 2.9 \text{ eV} \\ E_g = E_i &= 0.66 \text{ eV} \\ E_x &= 21 \text{ meV} \end{aligned}$$

$$\sigma_i = \frac{E_x}{E_i} \sqrt{\frac{E_0}{E_x} - \frac{E_i}{E_x} N_i} = \sqrt{\frac{E_0}{\epsilon_i}} \sqrt{\frac{E_x}{E_i} \left(\frac{\epsilon_i}{E_i} - 1 \right)} = \sqrt{F N_Q}$$





- Introduction to Germanium detectors
 - radiation detection in semiconductors
 - properties of Germanium detectors

- **Experimental setup**

- background measurement
- weak sources
- background reduction

- Data analysis

- calibration with KCl salt
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- what can we measure with XtRa?

Photons produced the decay chains of radioisotopes

Two kinds of measurement

Background measurements

=

everything else but a source



NATURAL RADIOACTIVITY

- cosmic radiation
- **terrestrial sources**
 - potassium
 - carbon
 - uranium and thorium
(decay chain elements)
- **human produced source**
 - Cs from nuclear explosions

Source measurements

1) EASY

- **strong sources**
- not present in natural background

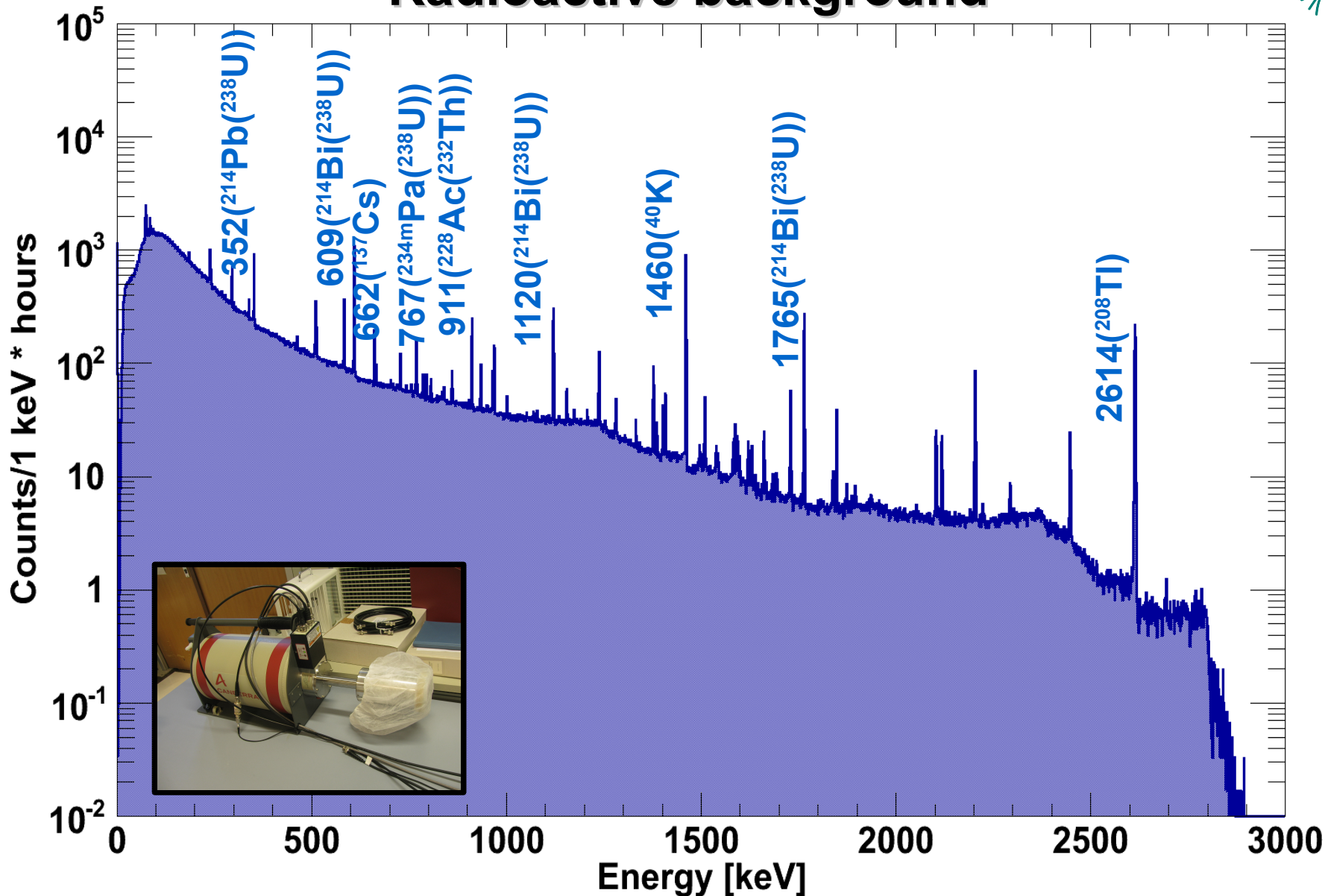
2) DIFFICULT

- **weak sources**
- present in natural background

**Energy spectra → IDENTIFY RADIOISOTOPES
→ QUANTIFY THEIR ABUNDANCE**



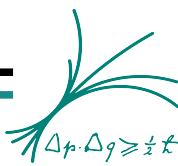
Radioactive background



- cannot measure a source without background
- **bkg SUBTRACTION is needed...** but is it good **enough** for weak sources?



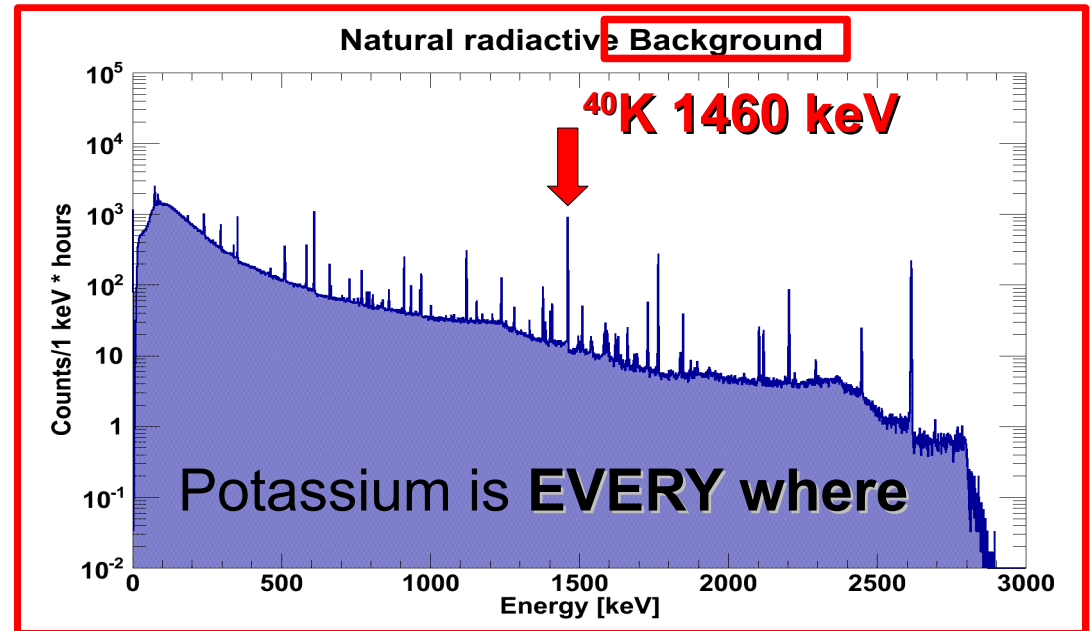
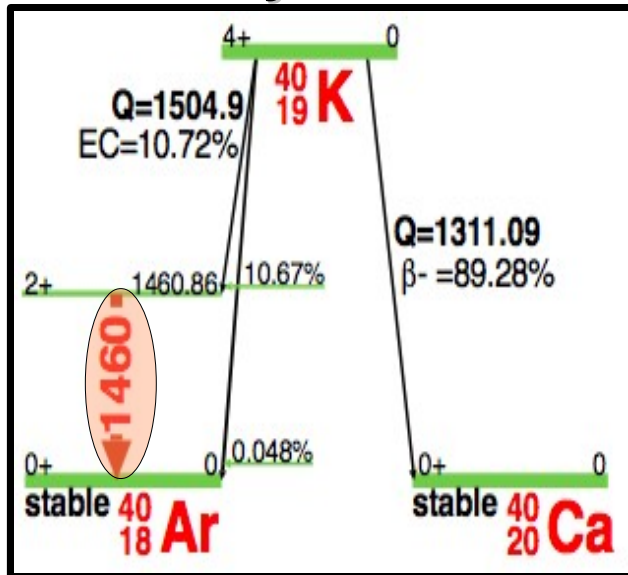
Experimental setup



Weak Sources: Potassium in food

- each food sample contains Potassium
- **100 g of strawberry** contains **153 mg of K** and **0.018 mg** of ^{40}K
 - small quantity of ^{40}K → small **activity** expected!!

Decay scheme



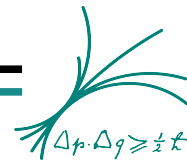
expected counts from the weak source in 1 hour → 20 counts
expected counts from the bkg radiation in 1 hour → 1000 counts



Signal is less than a fluctuation of the bkg!!! **WE CANNOT SEE IT!!**



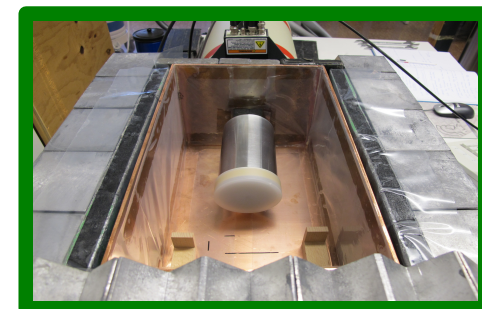
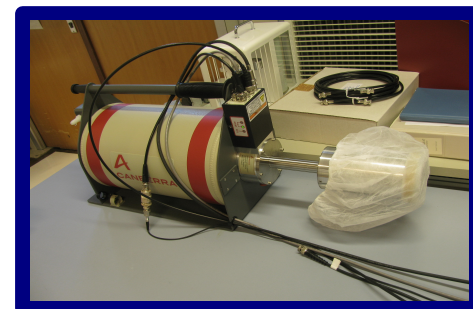
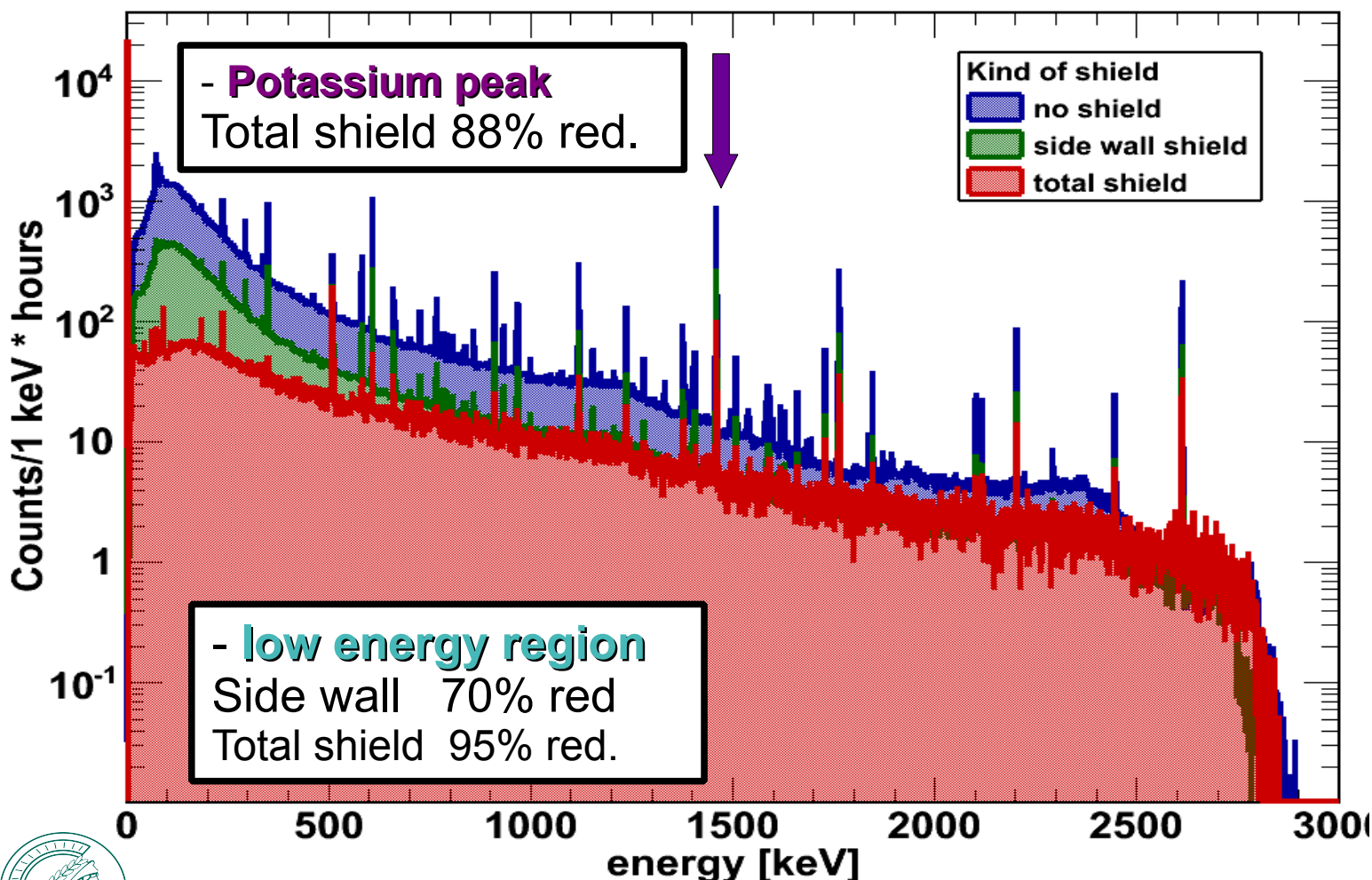
Experimental setup



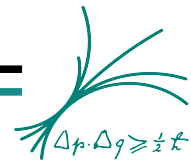
How can we **reduce the counts of background**?
- shielding the detector from natural radiation

LEAD SHIELD → high density and high Z can reduce the bkg!

Background reduction due to the lead castle

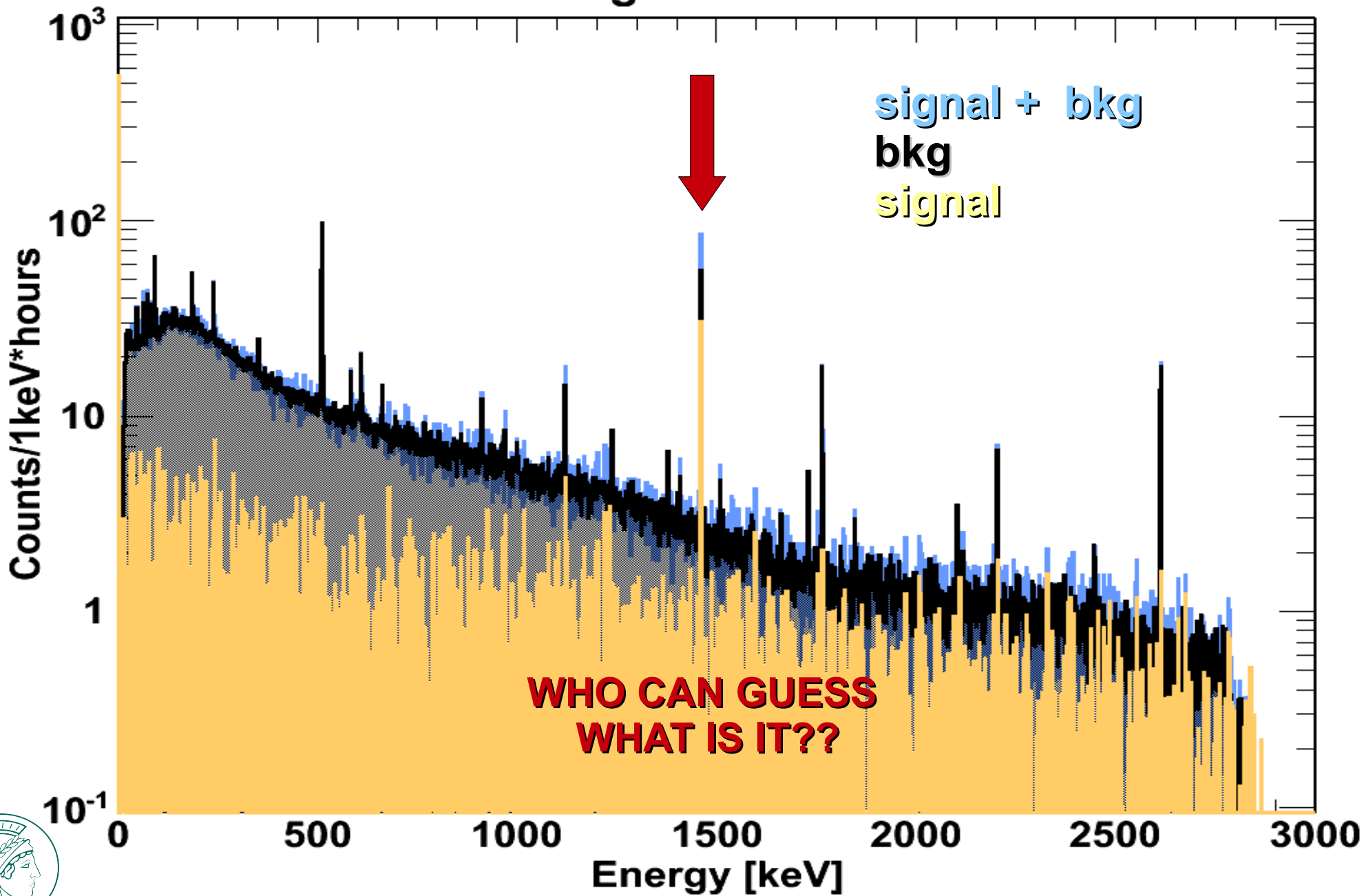


Experimental setup

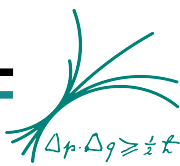


... and now... **YES WE CAN!!**

Effect of background subtraction

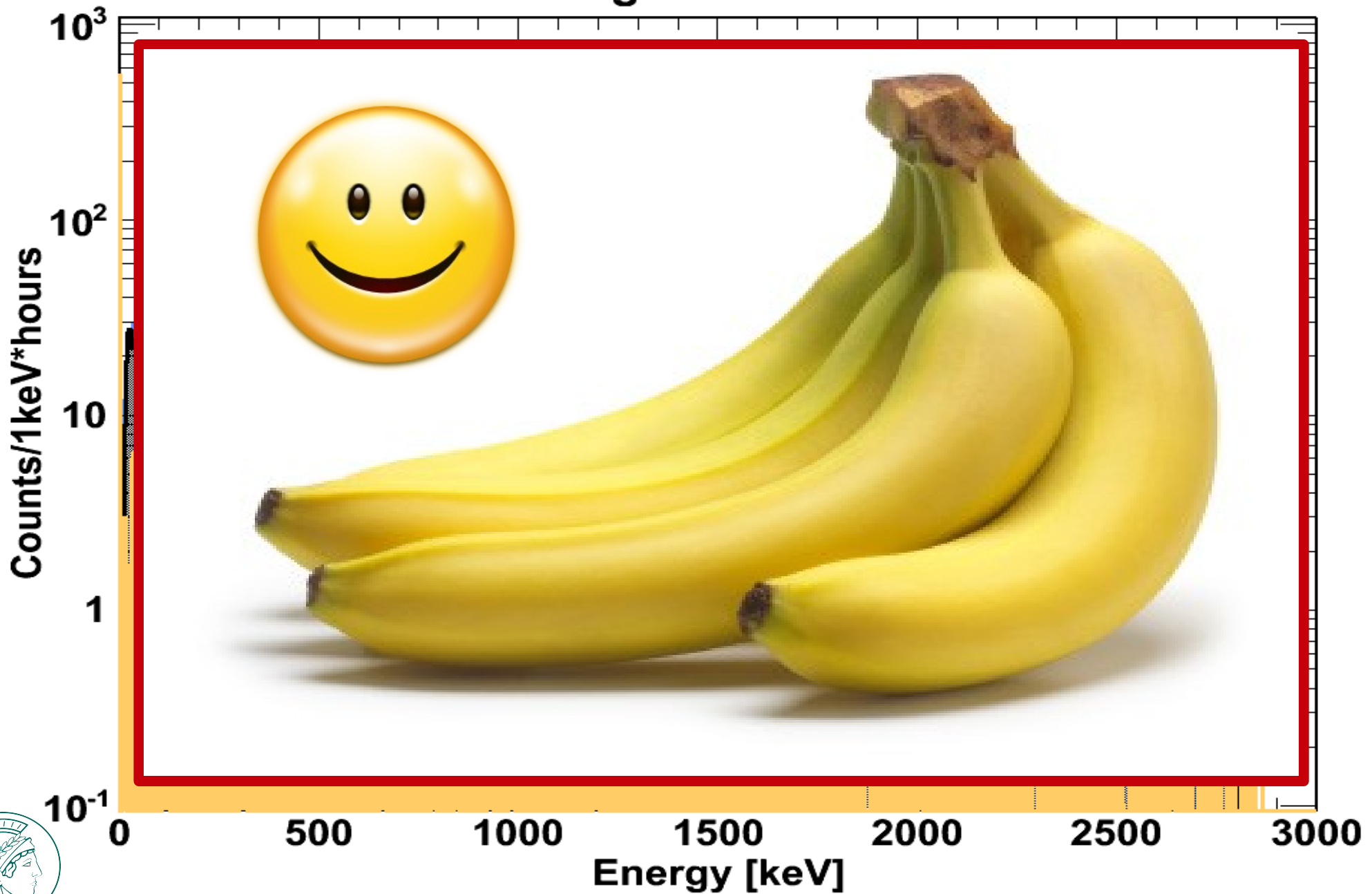


Experimental setup



... and now... **YES WE CAN!!**

Effect of background subtraction



→ **Introduction to Germanium detectors**

- radiation detection in semiconductors
- properties of Germanium detectors

→ **Experimental setup**

- bkg measurement
- strong and weak sources
- background reduction

→ **Data analysis**

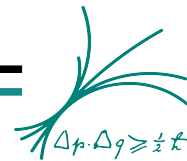
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- measurement of food rich in Potassium

→ **Conclusions**

- summary
- outlook



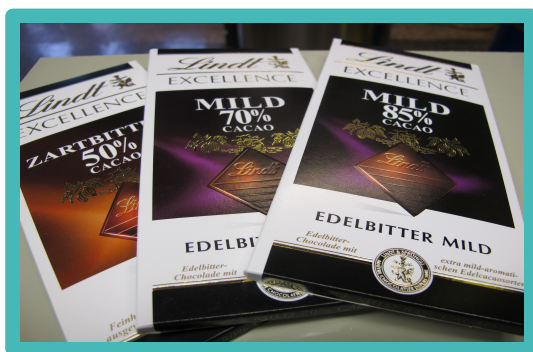
Data analysis



- with this experimental setup we can measure also **WEAK SOURCES!**
 - we can measure the **Potassium content** in different food sample

This are values that you can find in literature... but are they true??

Food	Potassium content in 1 pound [g]
Strawberry	0.76
Sugared almonds	1.27
White chocolate	1.43
kiwi	1.56
Banana	2.18
Hazelnuts	2.5
Dry Prunes	3.4
Raisins	3.74
Pistachios	5.125
Dry Apricots	9.25



Chocolate	Potassium content in 1 pound [g]
Dark Chocolate 50% cacao	3.81
Dark Chocolate 70% cacao	5.334
Dark Chocolate 85% cacao	6.477

We can do it but before...we need to **CALIBRATE** the detector!



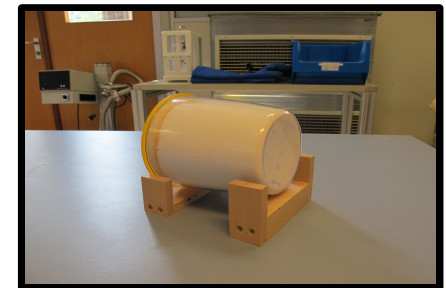
Check the **detector response** to a **WELL KNOWN** quantity of K

Known value of K : Spectrum_A = unknown value of K : Spectrum_B

Define the **experimental settings**:

1) **position of the plastic container:**

- vertical
- horizontal



2) **relative distance between source and detector**

MAXIMIZING THE GEOMETRICAL ACCEPTANCE

$$a^g = \frac{N_{measured}}{N_{expected}}$$

Where: $N_{measured}$ are obtained directly from the measured spectrum
 $N_{expected}$ can be **calculated from the activity** of the sample



Activity = number of decays per second

Directly from the exponential decay law for N_0 radioisotopes:

$$A = \frac{D(t)}{t} = N_0 \cdot \frac{\ln 2}{t_{\frac{1}{2}}}$$

if we have some amount m_I of radioisotope it will be:

$$A = \frac{D(t)}{t} = \left(\frac{m_I}{m_I^A} \cdot N_A \right) \cdot \frac{\ln 2}{t_{\frac{1}{2}}}$$

Potassium Chloride calibration salt

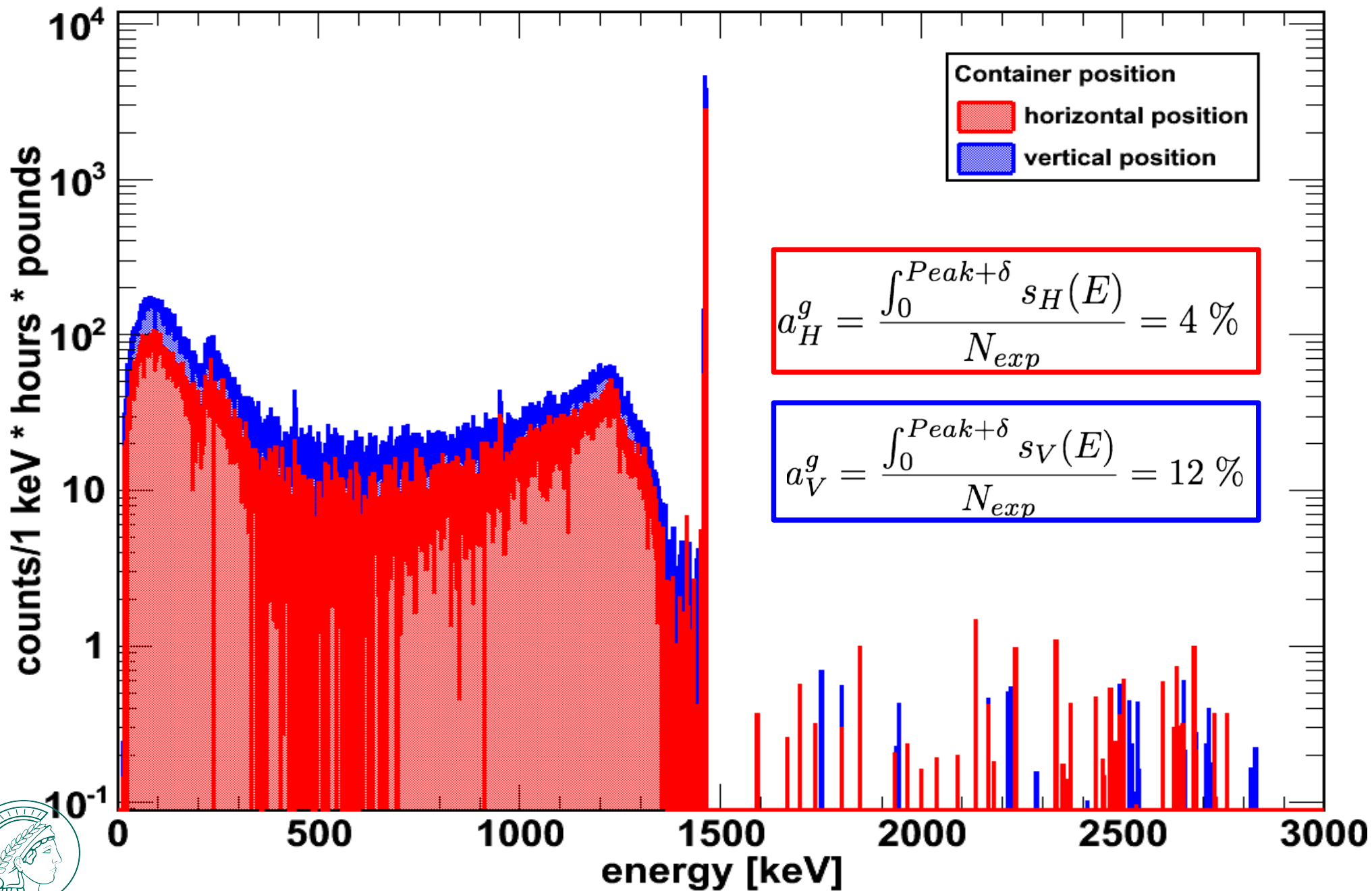
- **mass of the salt** sample
- **impurities**
- **mass of Potassium in KCl**
- **isotopic abundance** for ^{40}K
- **atomic mass of ^{40}K**

$$m_{^{40}\text{K}} = m_{\text{KCl}} \cdot (1 - i_{\text{tot}}) \cdot m_{\text{K}}^{\text{KCl}} \cdot a_{^{40}\text{K}}$$

$$A_{1460} = 141.9 \text{ Bq}$$



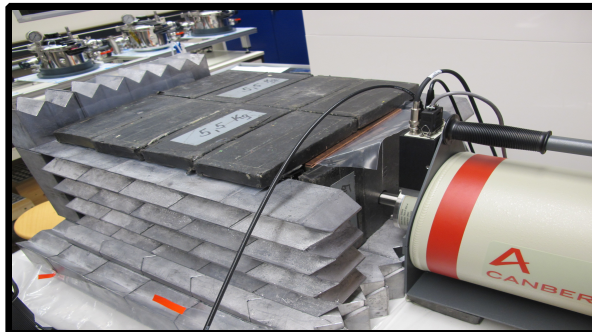
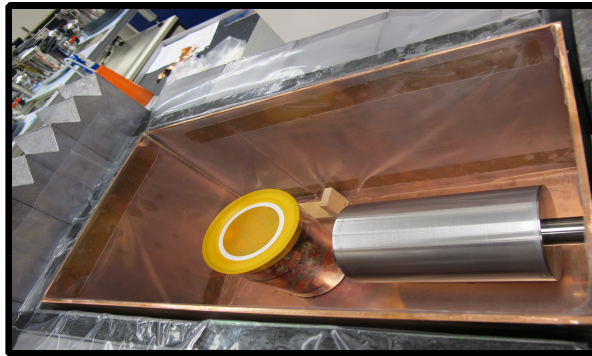
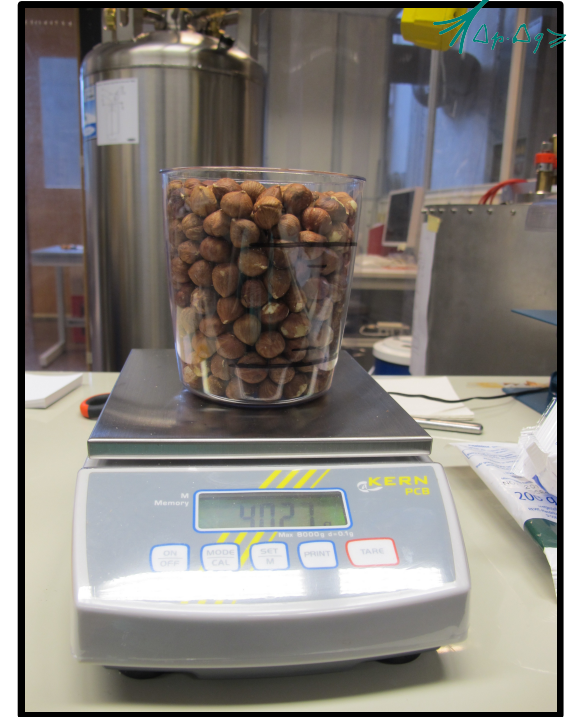
KCl spectra for different container positions



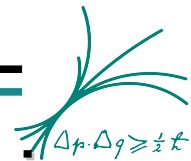
Data analysis



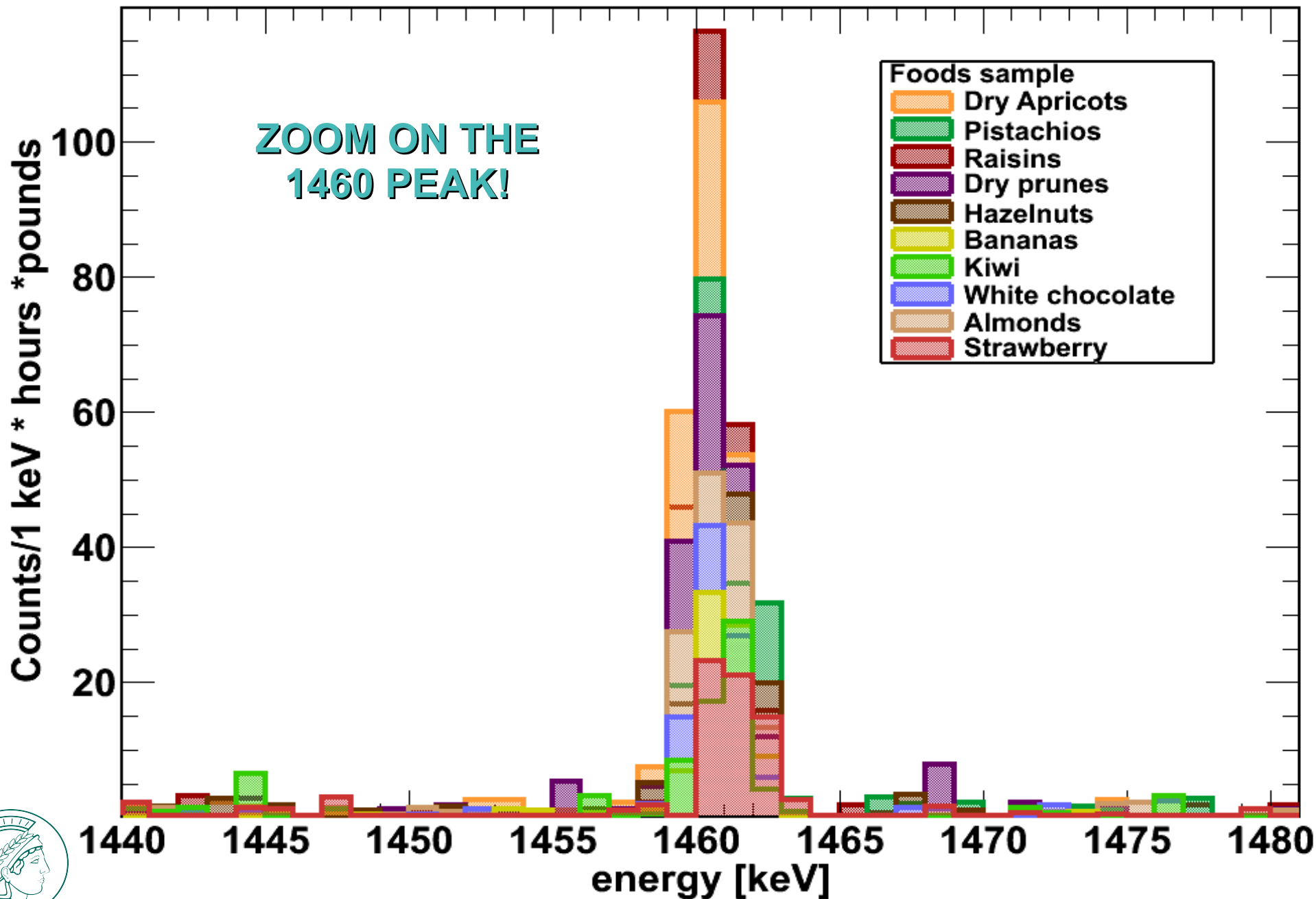
- all food samples **weighed**
- prepared in the **same plastic container**
 - try to have the **same acceptance**
- **self absorption neglected**



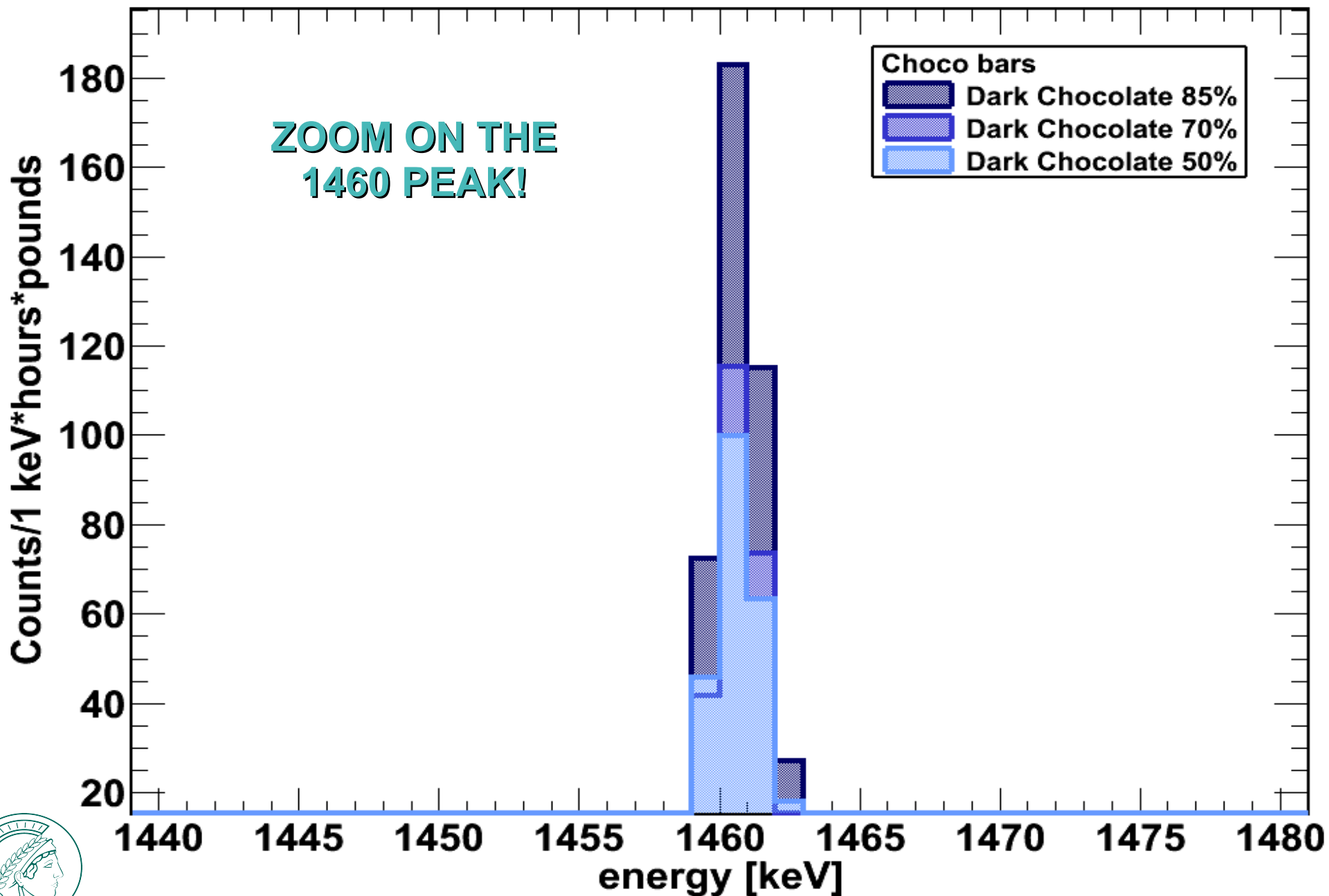
- plastic container always put **vertical**
- **same relative position of the source and the detector**



Comparison of spectra from different food sample rich in potassium



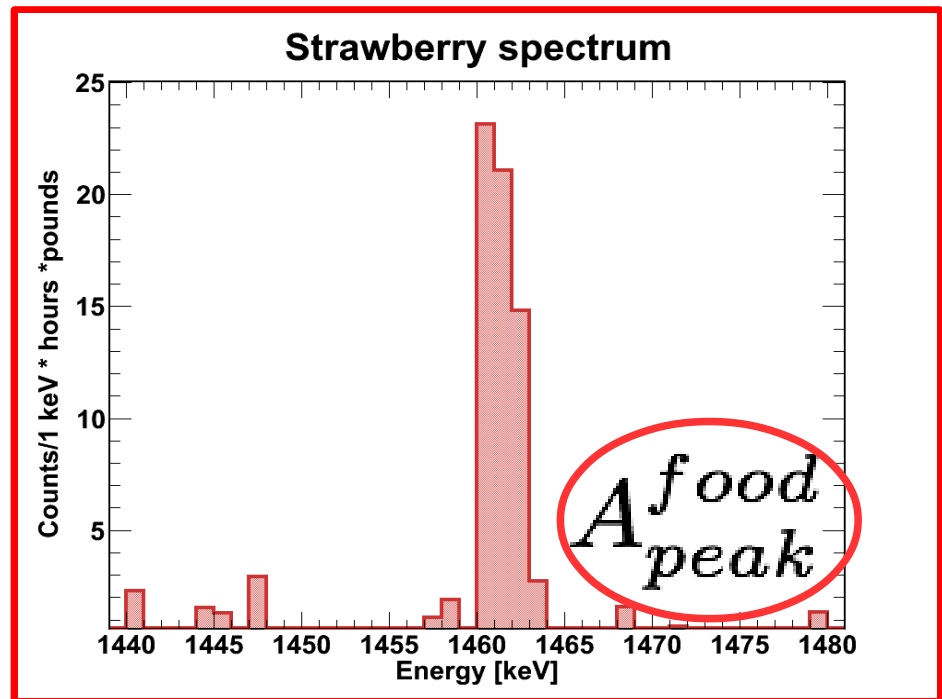
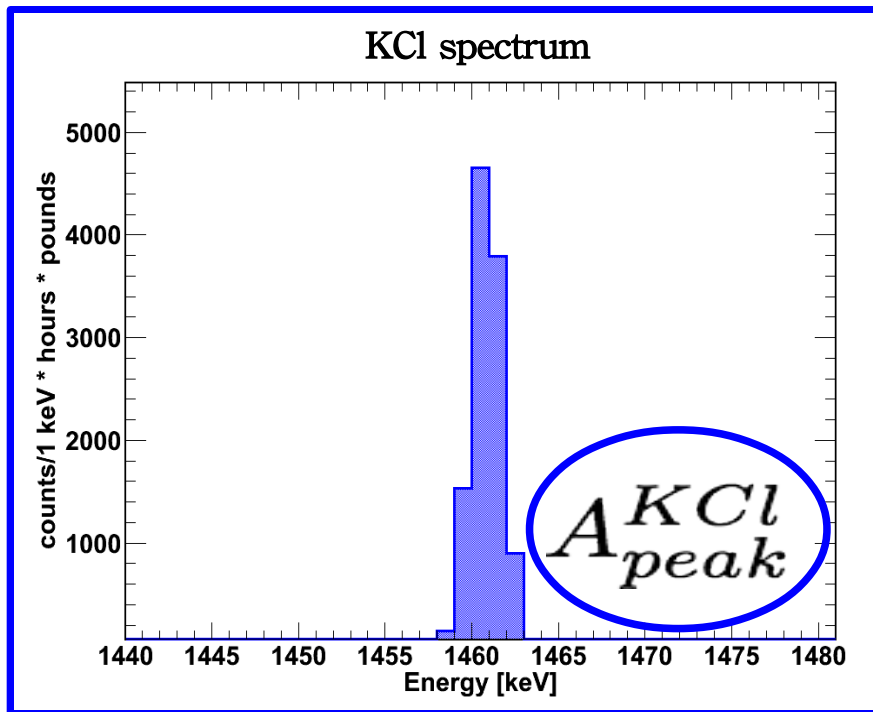
Comparison of spectra from different chocolate bars



To be more quantitative:

- we can use the already known proportion!!

Known value of K : Spectrum_A = unknown value of K : Spectrum_B

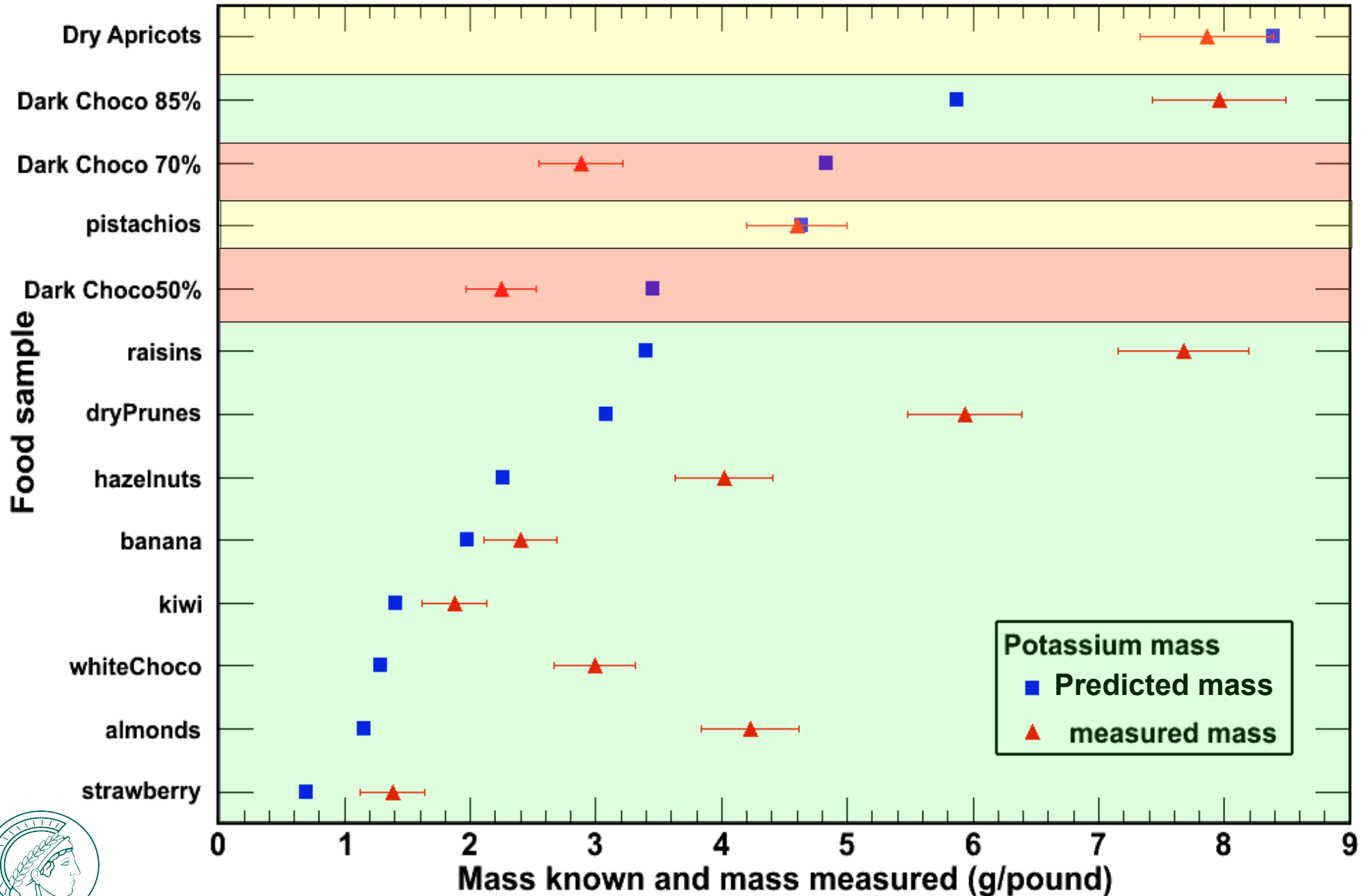


$$m_K^{meas} = \frac{A_{peak}^{food}}{A_{peak}^{KCl}} \cdot m_k^{KCl}$$





Potassium mass in food sample: known and measured value comparison



Just to summarize...

- built a shield with **high bkg reduction power**
- **good performances** of the detector with **weak sources** like strawberries
- measured **Potassium content** of **13** different food samples
- figured out a **non linear behaviour** of the **Potassium** content in different cacao percentages choco bars
 - Do they put **less cacao**?
 - Do they use **different kind of cacao**?



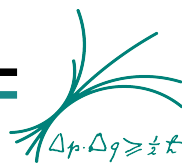
Outlook:

- **simulation** will be done to take into account **self absorption**



- **maybe...** **choco bars** measurement will be repeated with different **chocolate brands**

**BACK UP
SLIDES**



- **Signal formation** can be described with **events**:
 - with **low probability**
 - **independent** one from each other
 - with an **average rate** which doesn't change in the period of interest

EXPECTED VARIANCE = POISSON VARIANCE

OBSERVED VARIANCE = F * POISSON VARIANCE

with $F < 1$

Physical reason: ENERGY and MOMENTUM CONSERVATION

$$\begin{aligned} \epsilon_i &= 2.9 \text{ eV} \\ E_g = E_i &= 0.66 \text{ eV} \end{aligned}$$



- ionization \Rightarrow **electron-hole pairs**
- lattice excitation \Rightarrow **phonons**

$$E_0 = E_i N_i + E_x N_x$$

$$dE_0 = \frac{\partial E_0}{\partial N_x} dN_x + \frac{\partial E_0}{\partial N_i} dN_i = 0 \quad \Rightarrow \quad E_x \sigma_x = E_i \sigma_i$$

$$\sigma_i = \frac{E_x}{E_i} \sqrt{\frac{E_0}{E_x} - \frac{E_i}{E_x} N_i} = \sqrt{\frac{E_0}{\epsilon_i}} \sqrt{\frac{E_x}{E_i} \left(\frac{\epsilon_i}{E_i} - 1 \right)} = \sqrt{F N_Q}$$



Activity = number of decays per second

- from the **exponential decay law**: if we have initially N_0 radioisotopes

$$D(t) = N_0 - N(t) = N_0(1 - e^{-\frac{t}{\tau}})$$

is the number of decays after a time t

- if we use the half life $t_{1/2}$ and expand in **Taylor series** we can obtain

$$e^{-\frac{t}{\tau}} = 1 - \frac{t}{\tau} = 1 - \frac{t \cdot \ln 2}{t_{1/2}}$$

- if we have a **certain mass of a radioactive isotope** m_I

$$A = \frac{D(t)}{t} = \left(\frac{m_I}{m_I^A} \cdot N_A \right) \cdot \frac{\ln 2}{t_{1/2}}$$

For a Potassium Chloride (KCl) sample :

- of mass m_{KCl}
- with a certain amount of impurities i_{tot} we will have:

$$m_{40K} = m_{KCl} \cdot (1 - i_{tot}) \cdot m_K^{KCl} \cdot a_{40K}$$

$$A_{1460} = 141.9 \text{ Bq}$$

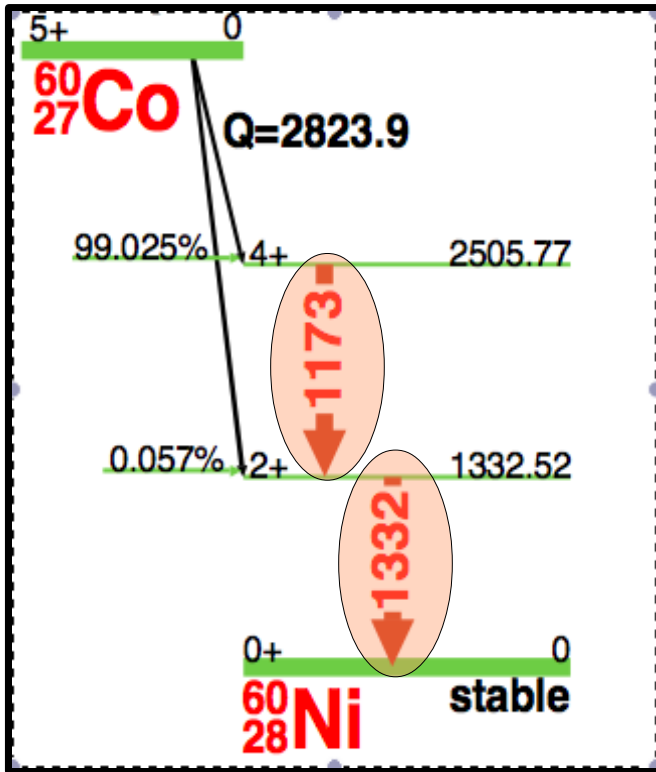


Experimental setup

Name: $^{228}_{90}\text{Th}$
 Neutrons: 138
 Activity:



Decay chain:

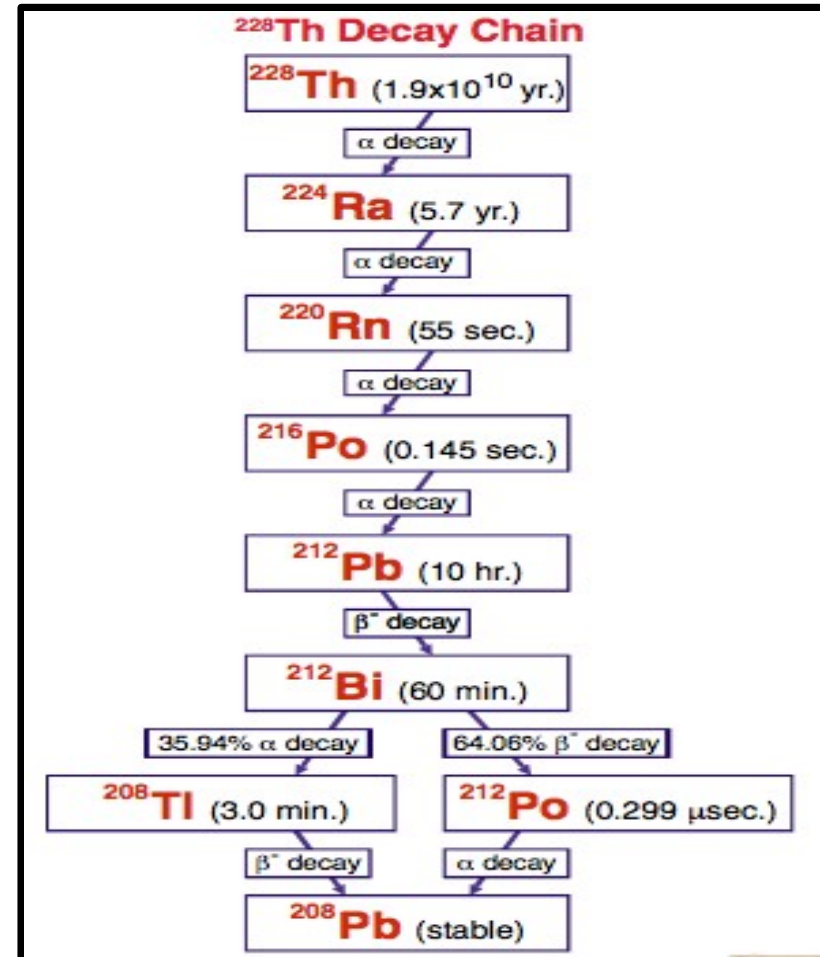


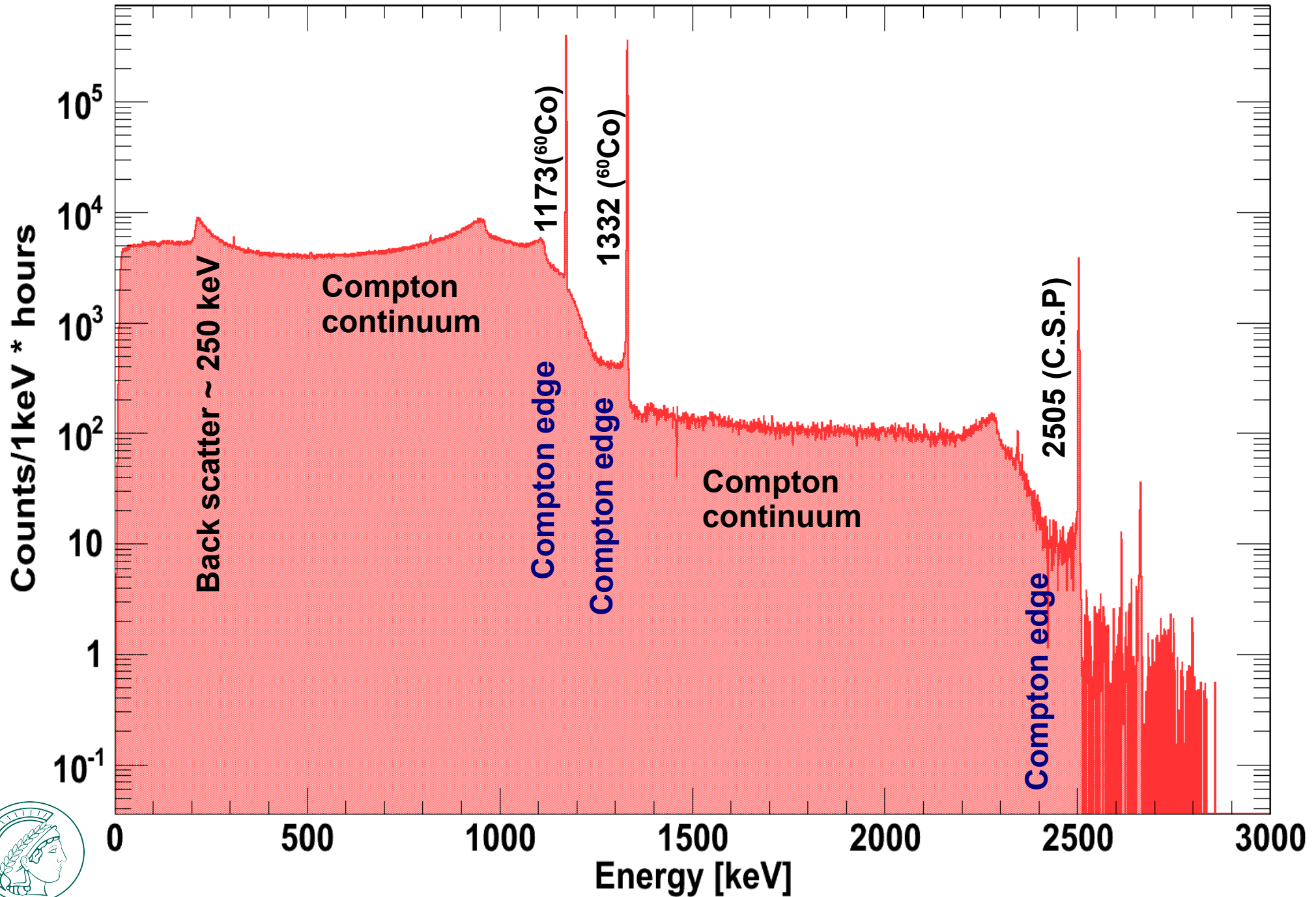
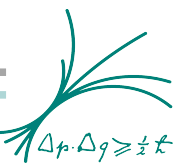
Isomeric transition with **photon emission**

Name: $^{60}_{27}\text{Th}$
 Neutrons: 33
 Activity:



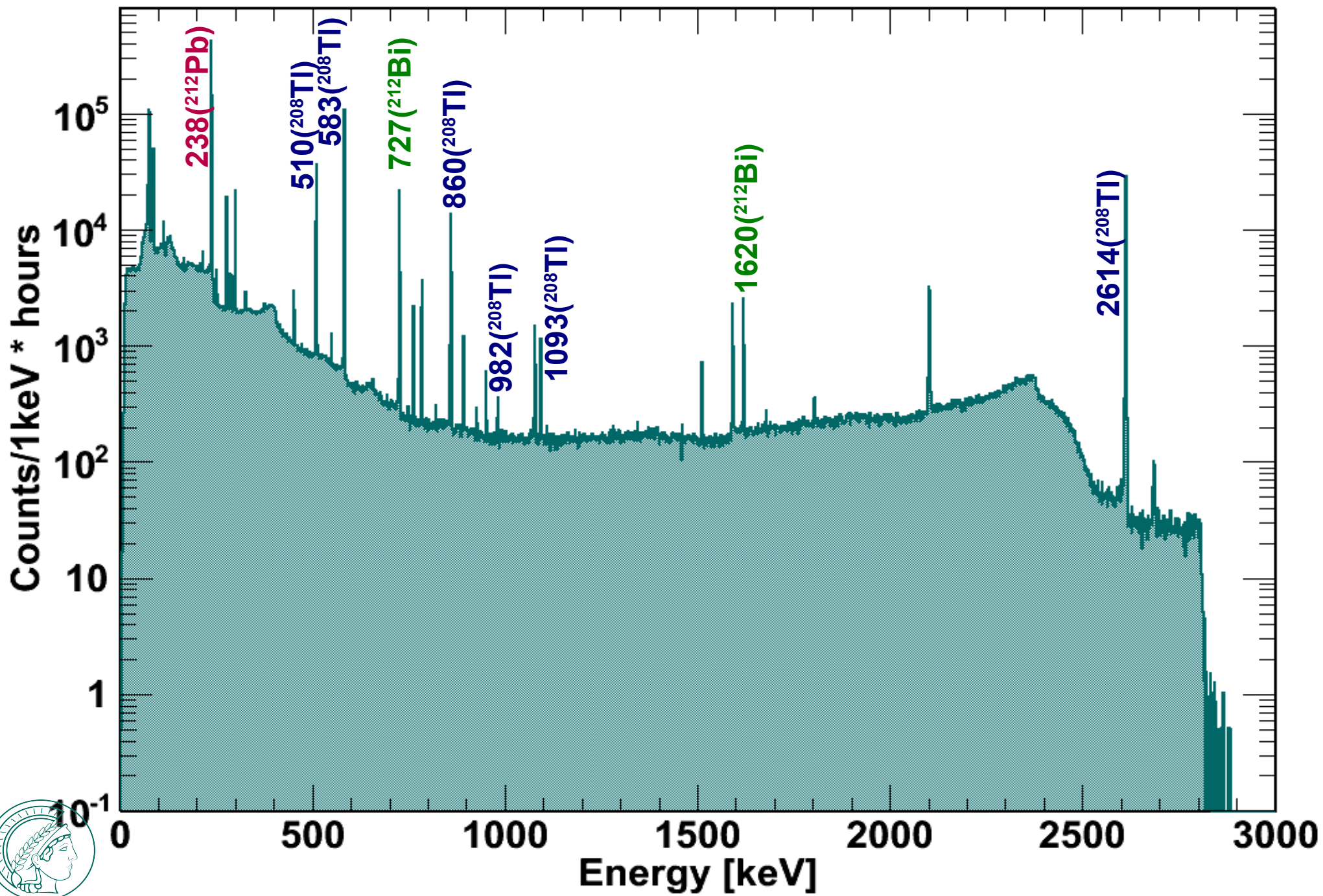
Decay chain:



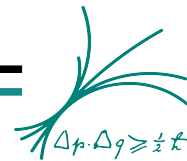




Th228 spectrum



Data analysis



We can use strong sources to know the detector....

Fano Factor Measurement

→ energy resolution depends on ENERGY

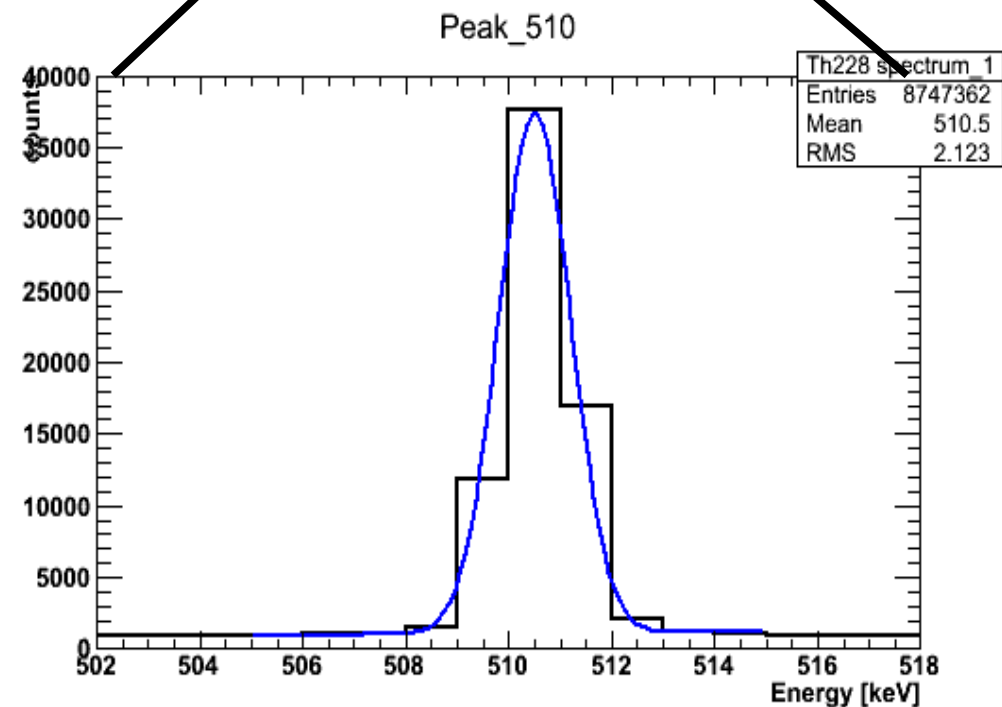
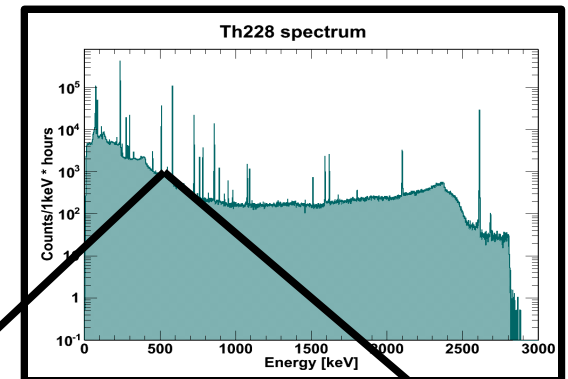
$$FWHD_{tot}^2 = (2.35)^2 F \epsilon E + b^2$$

→ scanning all the energy range in the ^{228}Th spectrum
- choosing good peaks

238 keV	^{212}Pb
510 keV	^{208}Tl
583 keV	^{208}Tl
727 keV	^{212}Bi
860 keV	^{208}Tl
982 keV	^{208}Tl
1093 keV	^{208}Tl
1620 keV	^{212}Bi
2614 keV	^{208}Tl

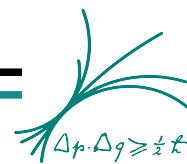
- fit with a Gaussian
=> extract the RMS

- check the dependence on energy



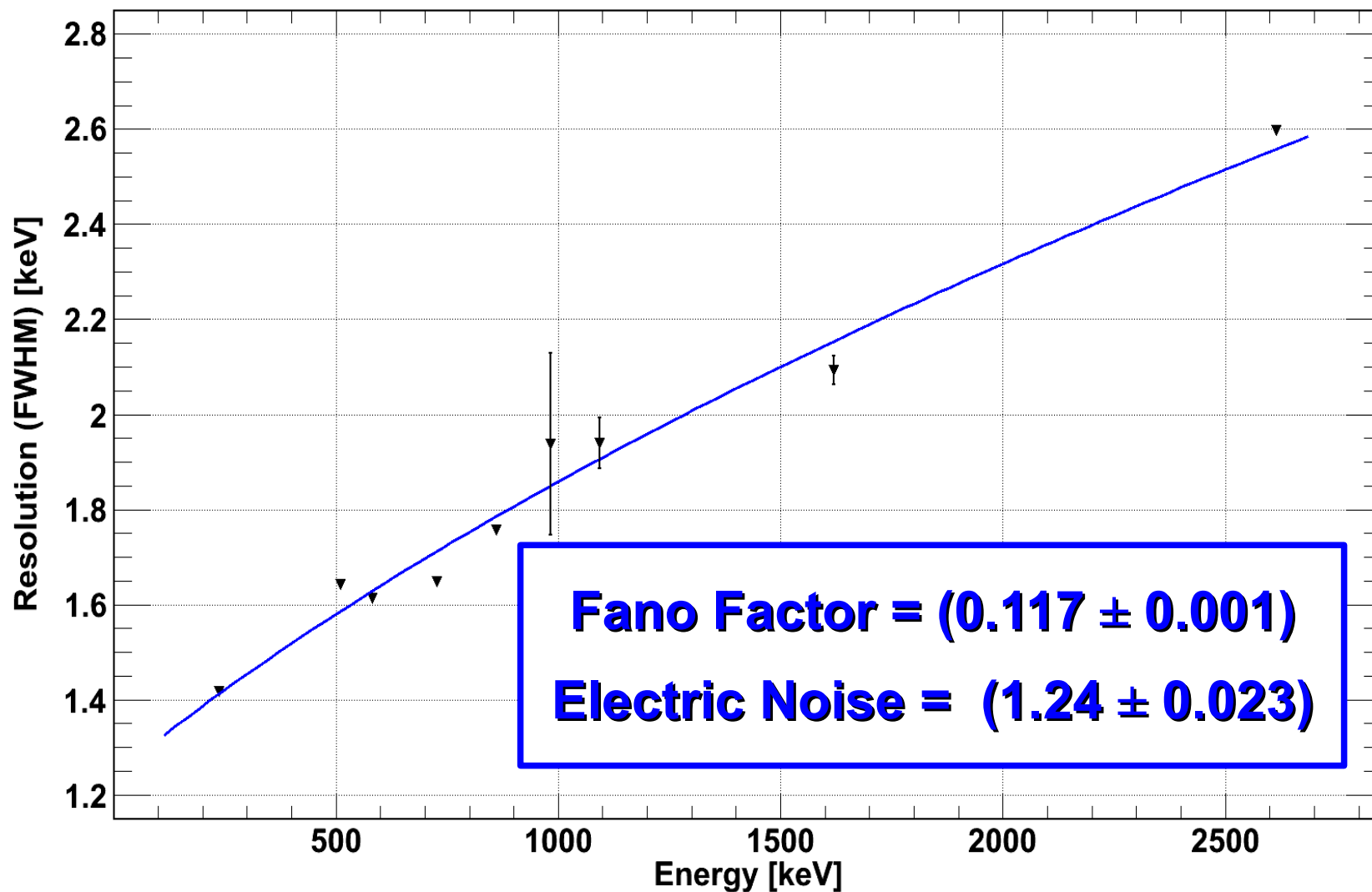
$$RMS = (0.699 \pm 0.0026) \text{ keV}$$





$$FWHM \pm \sigma_{FWHM} = 2.35 \cdot (RMS \pm \sigma_{RMS})$$

Energy dependence of the resolution



To be compared with the theoretical value: **F = 0.13 in Ge**





Th228 spectrum

