

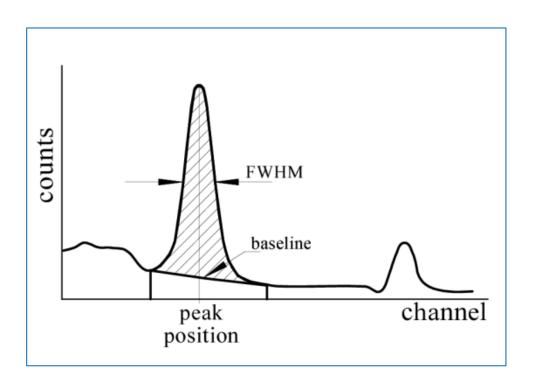
# Peak area determination in γ-ray spectrometry

Influence caused by subjective elements

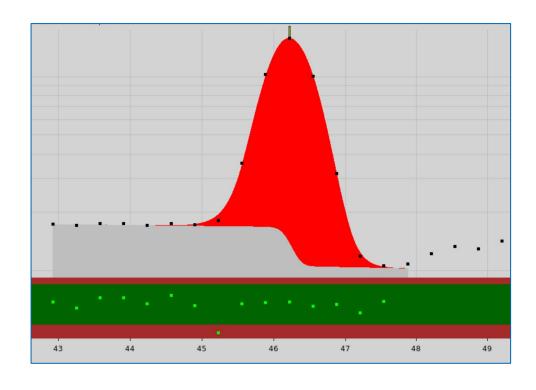
Roy Pöllänen

## 1. Two main approaches in the peak area determination

### **ROI-method:**



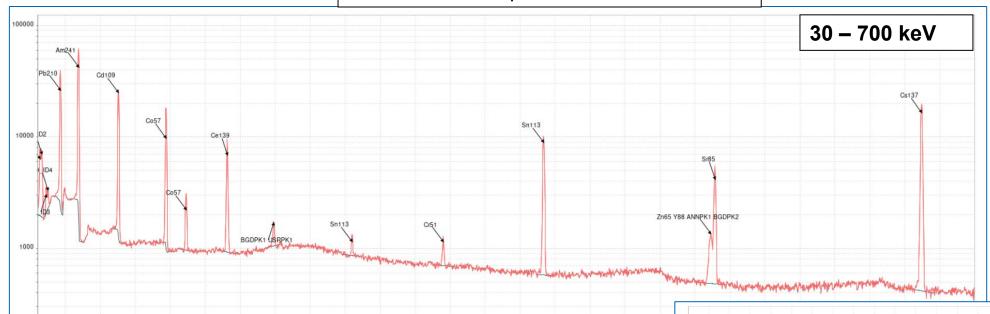
### **Peak fitting:**





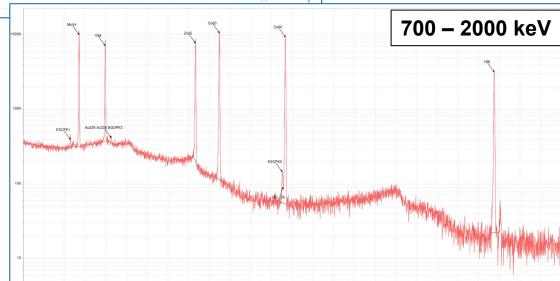
# 2. Mixed nuclide energy spectrum (for the efficiency calibration)

Several nuclides present: Pb-210 ... Y-88

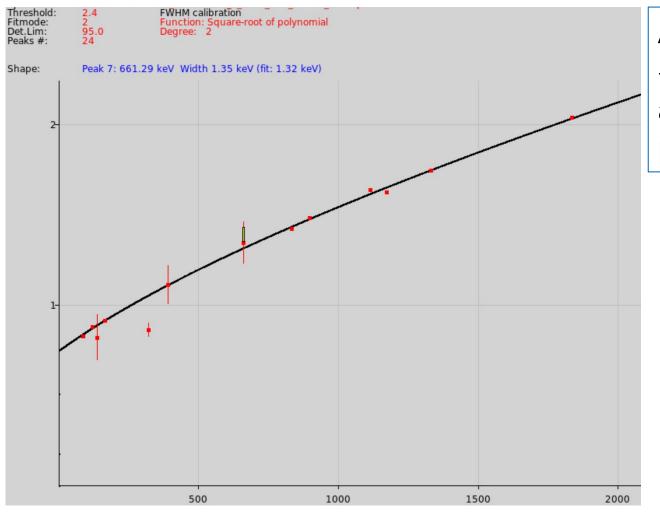


- Simple case from the point of view of peak area determination.
- But does the person analyzing the spectrum have an effect on the determined peak areas?



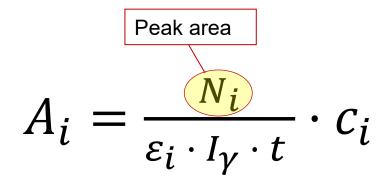


# 3. Example case: peak fitting by using Unisampo program



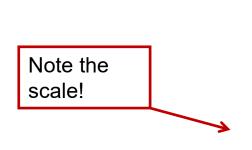
Automated peak shape calibration

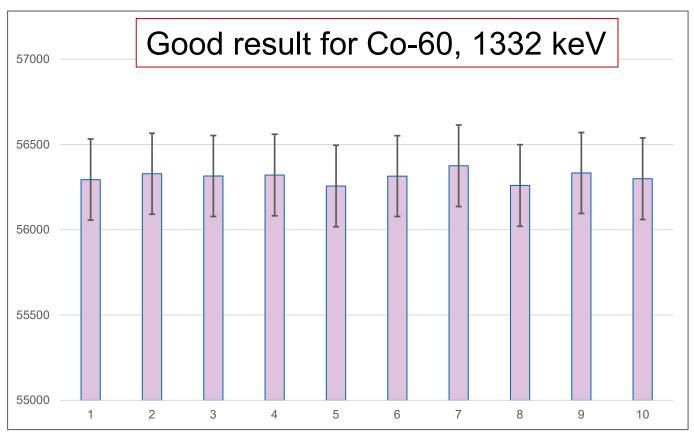
→ the analyst can tailor it! (as well as other parameters affecting the peak area)





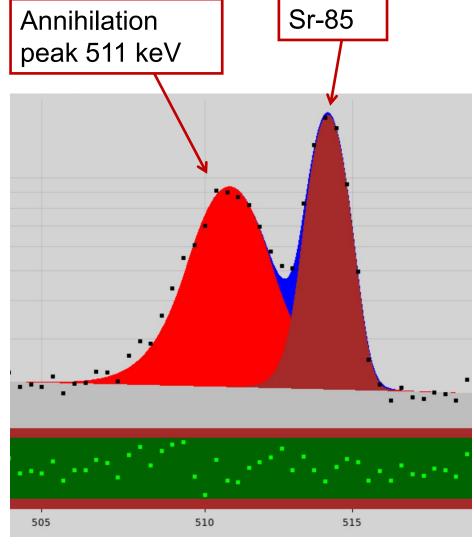
# 10 experienced gamma spectrum analyzers independently calculated peak areas from the <u>same spectrum</u>



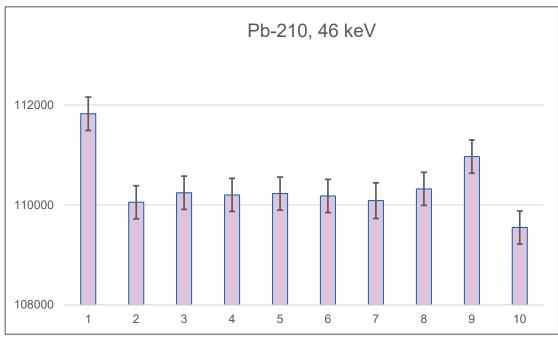


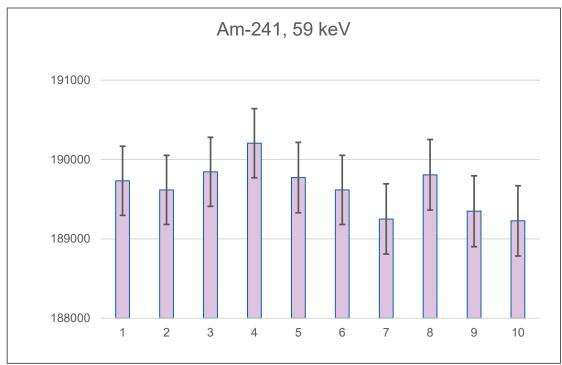


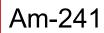


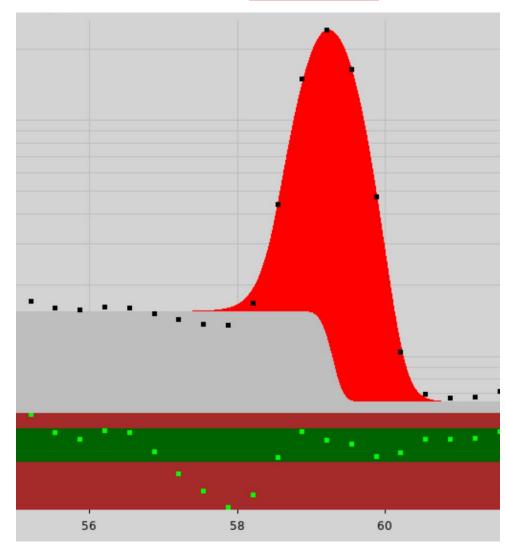




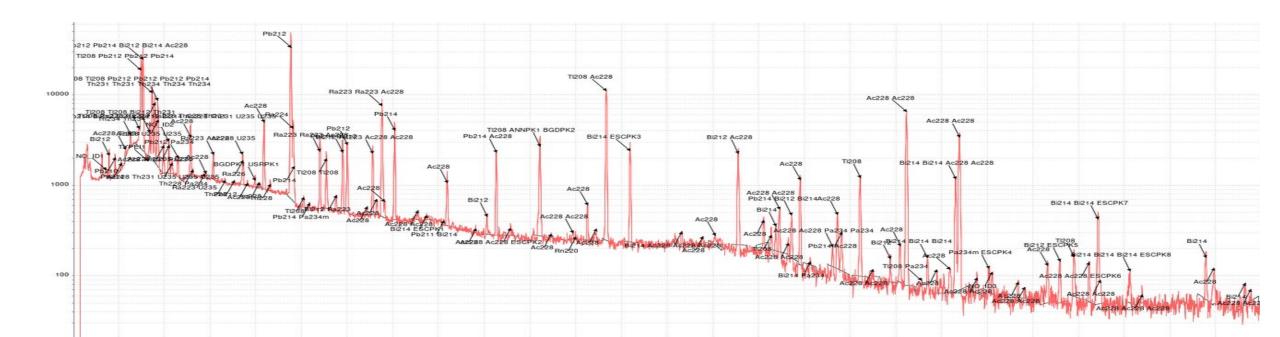








- Although the example spectrum was easy to analyze there were several points that had to be taken into account:
  - Using step function (or not)
  - Neighbouring peaks (such as Sr-85 and 511 keV annihilation peak)
  - Escape peaks (e.g. Co-60: 1173 keV 511 keV = 662 keV  $\rightarrow$  Cs-137: 661 keV)
  - Compton edges (e.g. Co-60 edge at the energy of around 1115 keV  $\rightarrow$  Zn-65: 1116 keV)
  - Lots of other things/effects ...
- What about then when a more complex spectrum must be analyzed?



### 4. Conclusion

Quality & metrological perspective:

How to take human-related factors into account especially in the uncertainty estimation?



# comments, please!

