

Hands-on session: Radioactivity Laboratory

Spectrometry with NaI(Tl)

Prof. Giuseppe Iaselli

Dott. Dayron Ramos

Dott. Nicola Ferrara



Istituto Nazionale di Fisica Nucleare

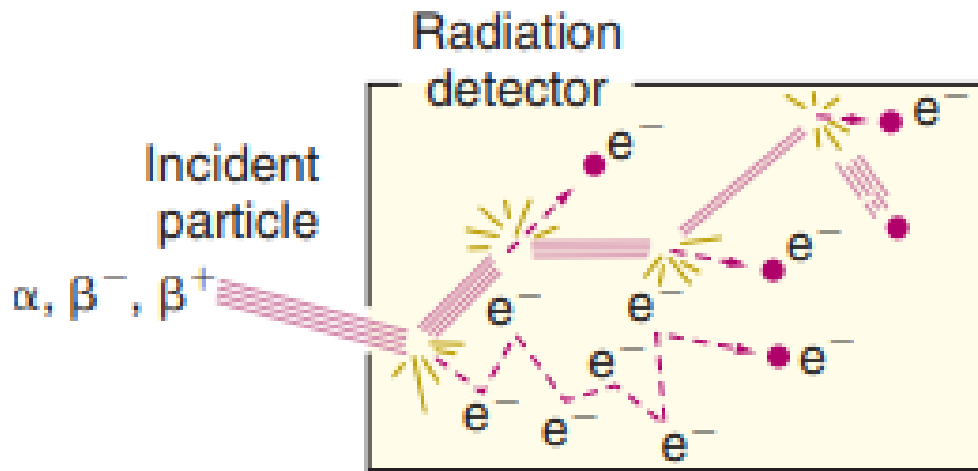


Politecnico
di Bari

Gamma Spectrometry

- Introduction to spectrometry
- NaI(Tl) photopeak detection
- Detection response
- Multi-Channel-Analysis
- Measurements

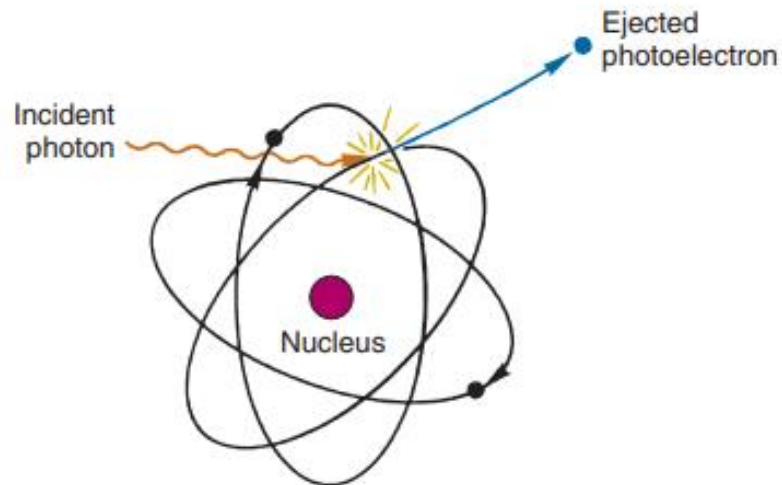
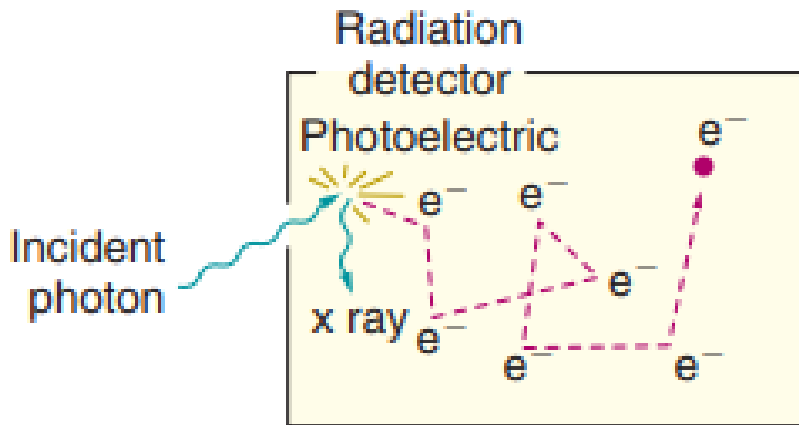
Basics of Pulse-Height Spectrometry



- Particles deposit energy into sensitive volume of detector
- Signal output proportional to particle energy
- Energy deposited outside the detector is not registered

Basics of Pulse-Height Spectrometry

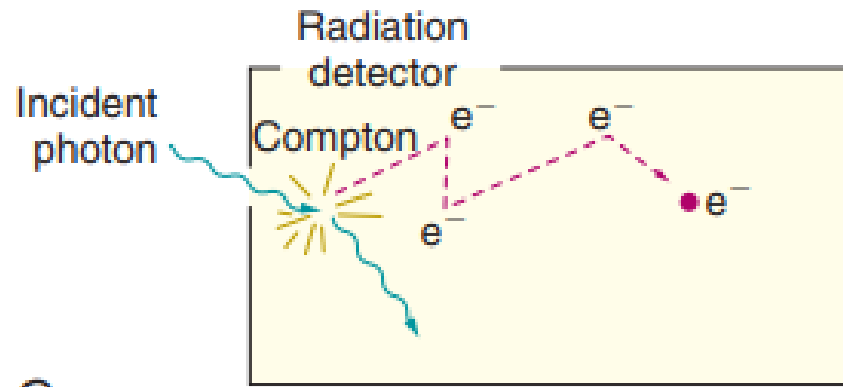
Photoelectric effect



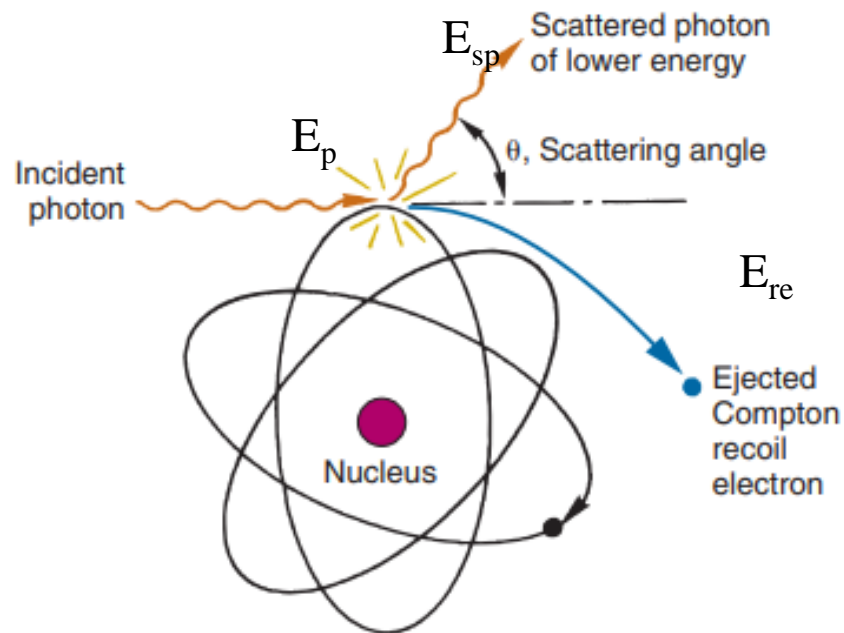
- Energy of photon transferred to atom
- Atom ejects an electron
- Photon is absorbed
- $E_{pe} = h\nu - K_B$

Photoelectron ejection creates a vacancy in an orbital electron shell, which is occupied by an electron of higher shell, which in turns lead to emission of a x-ray or an emission of an Auger electron

Basics of Pulse-Height Spectrometry



C



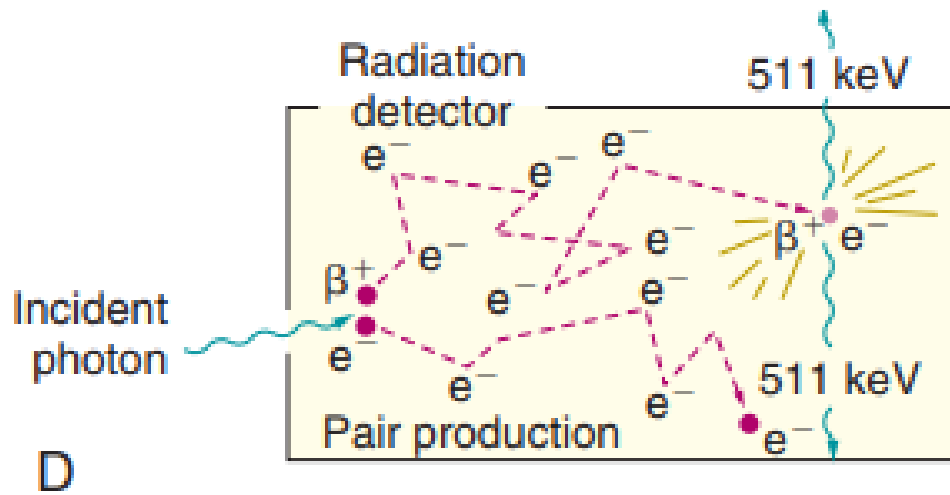
Compton effect:

- Energy of photon changes, because is transferred to electron
- Photon is not absorbed, but is deflected by θ angle

$$E_{sf} = \frac{E_f}{\left(1 + \frac{E_f}{0.511}(1 - \cos\theta)\right)}$$

- E_{sf} energy of scattered photon
- E_f energy of incident photon

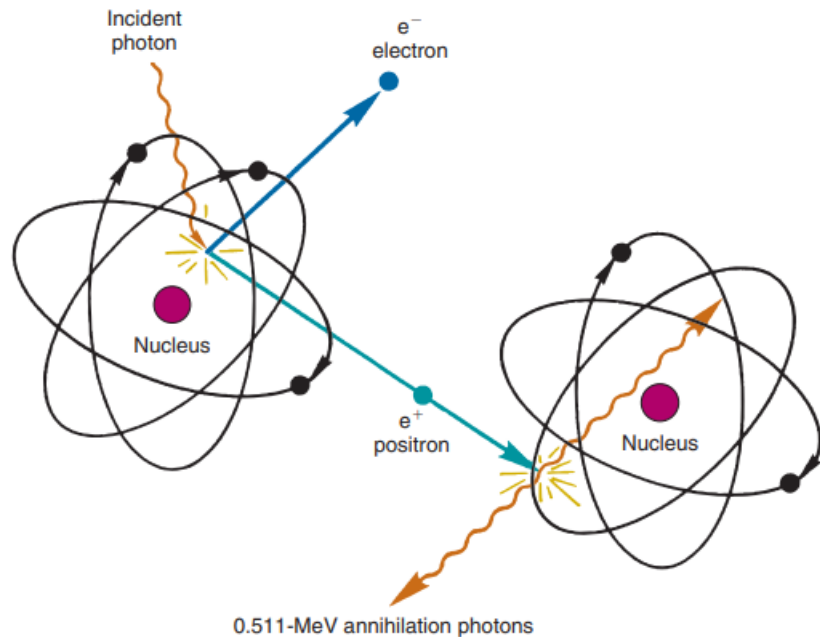
Basics of Pulse-Height Spectrometry



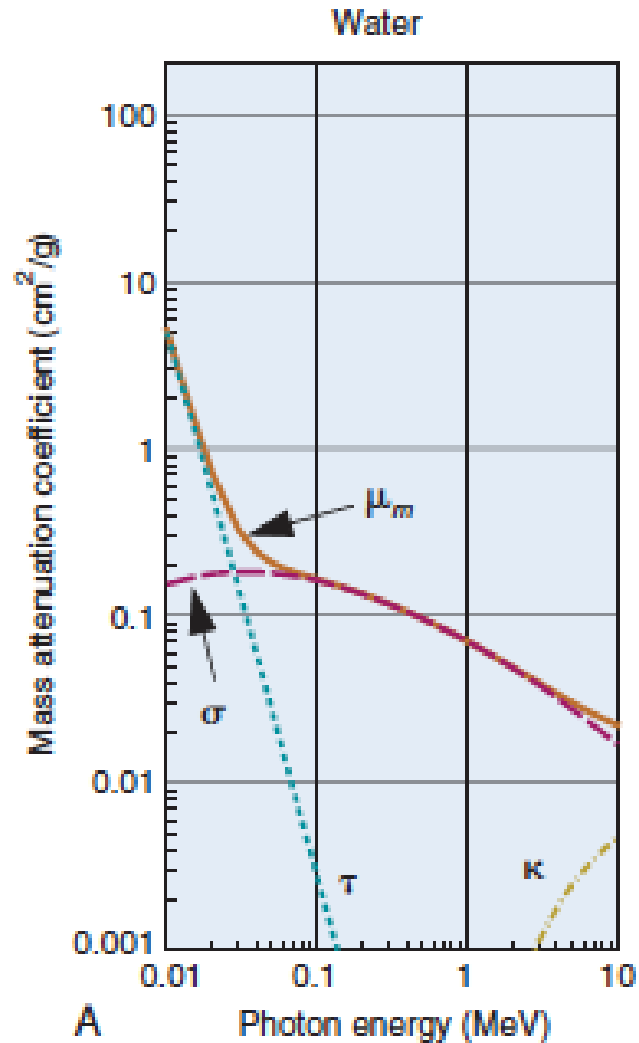
Pair production:

Photon undergoes pair production if energy is more than 1022 keV

Positron undergoes annihilation with an electron, producing two photons back-to-back



Basics of Pulse-Height Spectrometry



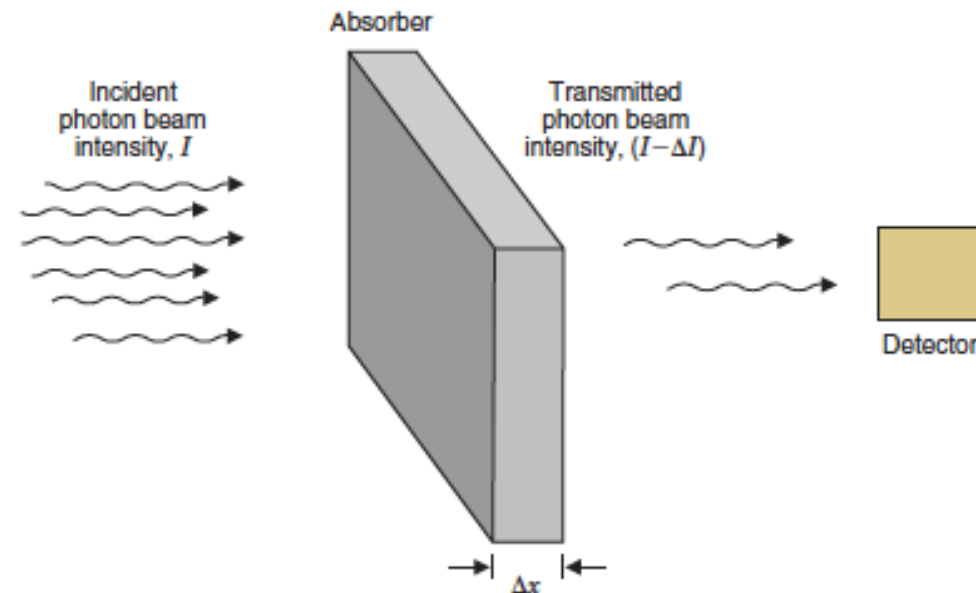
Attenuation coefficient:

$$N = N_0 e^{-\mu(E)x}$$

Mass attenuation coefficient:

$$\mu_m = \frac{\mu}{\rho}$$

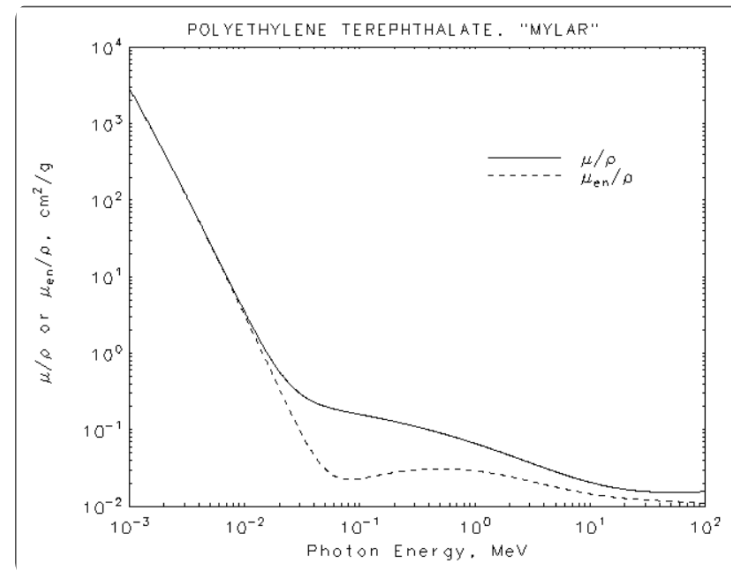
It depends on energy and on type of interaction



Energy (MeV)	μ/ρ (cm ² /g)	μ_{en}/ρ (cm ² /g)
1.00000E-03	2.911E+03	2.905E+03
1.50000E-03	9.536E+02	9.513E+02
2.00000E-03	4.206E+02	4.192E+02
3.00000E-03	1.288E+02	1.279E+02
4.00000E-03	5.466E+01	5.398E+01
5.00000E-03	2.792E+01	2.737E+01
6.00000E-03	1.608E+01	1.561E+01
8.00000E-03	6.750E+00	6.370E+00
1.00000E-02	3.481E+00	3.153E+00
1.50000E-02	1.132E+00	8.668E-01
2.00000E-02	5.798E-01	3.462E-01
3.00000E-02	3.009E-01	9.972E-02
4.00000E-02	2.304E-01	4.695E-02
5.00000E-02	2.020E-01	3.082E-02
6.00000E-02	1.868E-01	2.508E-02
8.00000E-02	1.695E-01	2.247E-02
1.00000E-01	1.586E-01	2.297E-02
1.50000E-01	1.406E-01	2.567E-02
2.00000E-01	1.282E-01	2.772E-02
3.00000E-01	1.111E-01	2.990E-02
4.00000E-01	9.947E-02	3.073E-02
5.00000E-01	9.079E-02	3.093E-02
6.00000E-01	8.395E-02	3.079E-02
8.00000E-01	7.372E-02	3.005E-02
1.00000E+00	6.628E-02	2.909E-02
1.25000E+00	5.927E-02	2.780E-02
1.50000E+00	5.395E-02	2.657E-02
2.00000E+00	4.630E-02	2.444E-02
3.00000E+00	3.715E-02	2.135E-02

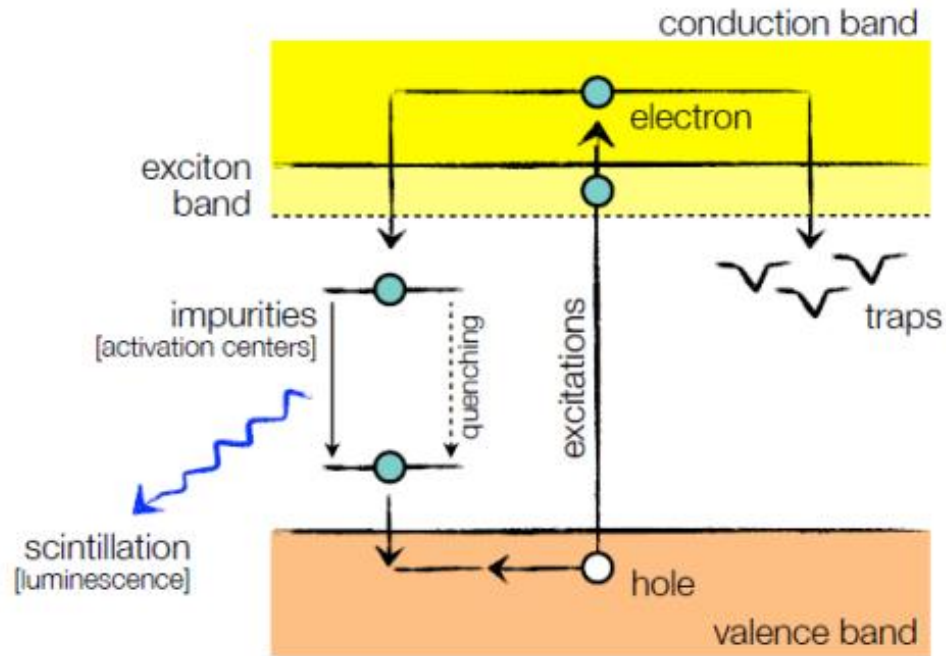
NIST database

- Table shows mass attenuation coefficient at different energy for the polyethylene terephthalate



- Density=1.38 g/cm³

Spectrometry with NaI(Tl)



Energy bands in impurity activated crystal showing excitation, luminescence, quenching and trapping

- Inorganic scintillators are lab grown crystals.
- Pure crystals make bad scintillators:
 - the defined energy bands between valence band and excitation band means that energy emitted by the crystal get reabsorbed by the same lattice
- There are only definite energy quanta allowing for the excitation of molecules
- Adding impurity improves the behavior
- There are more intermediate states between valence and excitation band that can be reached by the excited electron
- The emitted light is not completely reabsorbed by the crystal lattice

Spectrometry with NaI(Tl)

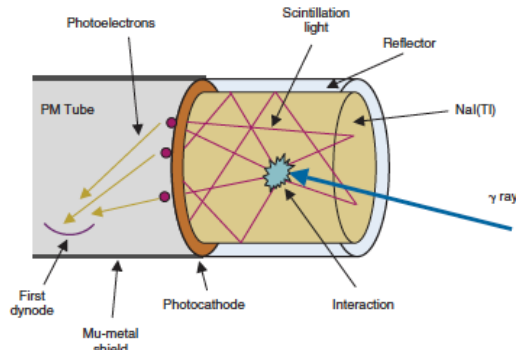


FIGURE 7-16 Arrangement of NaI(Tl) crystal and photomultiplier (PM) tube in a typical scintillation detector assembly.

- Good coupling with Photomultiplier by the use of light guide

- Multiplication of electrons, accelerated from a dynode to the other

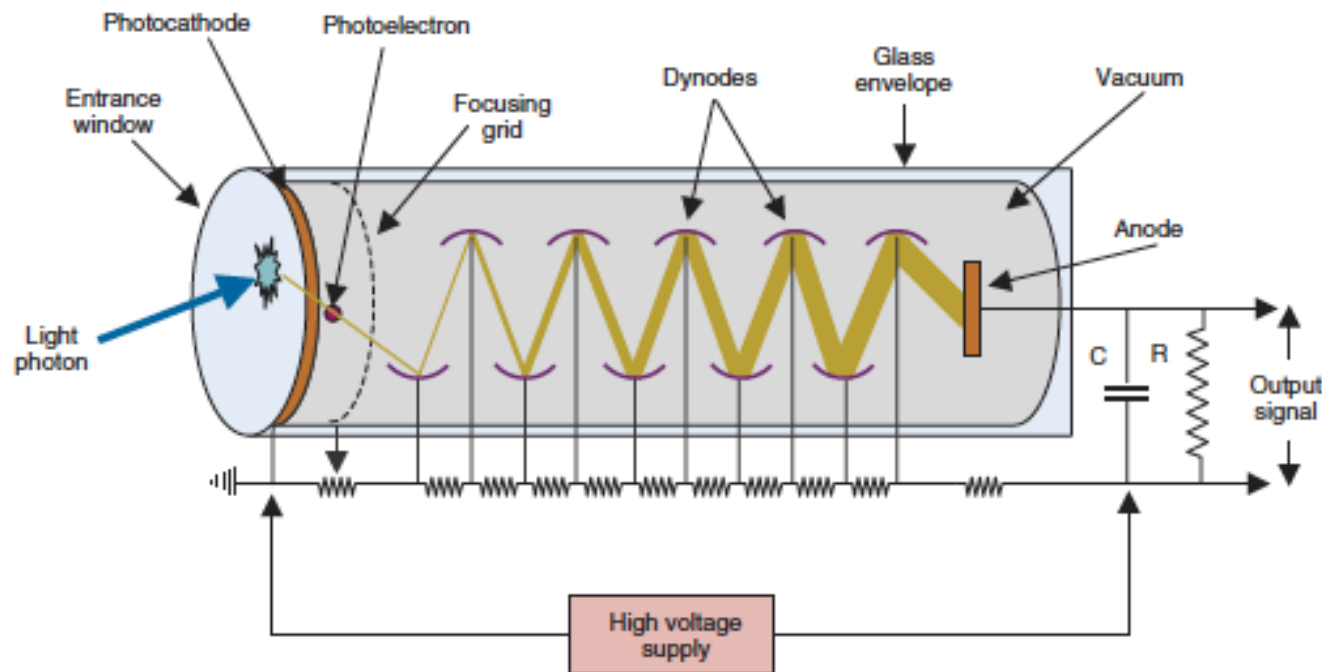
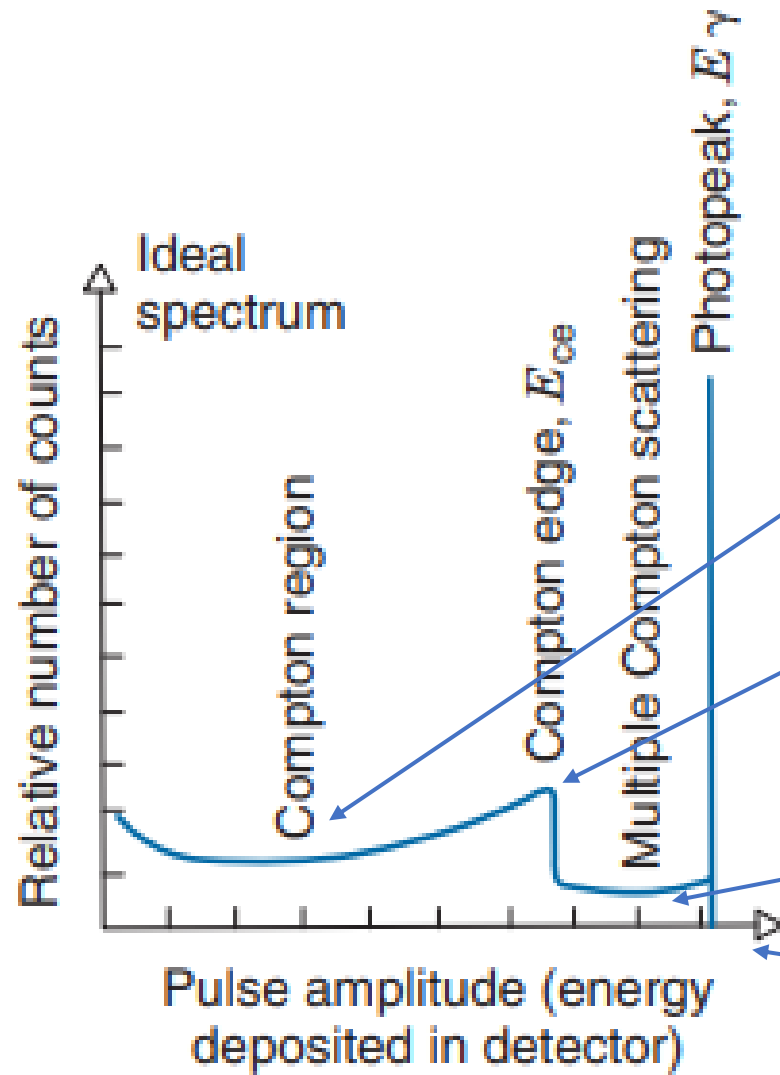


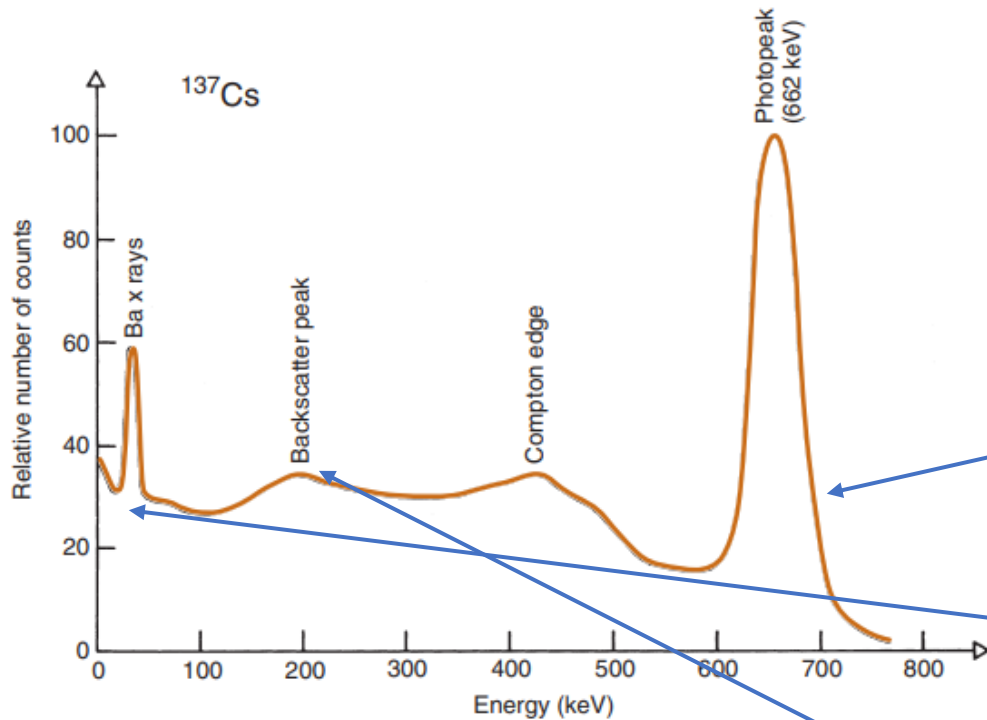
FIGURE 7-13 Basic principles of a photomultiplier tube.

Ideal spectrum



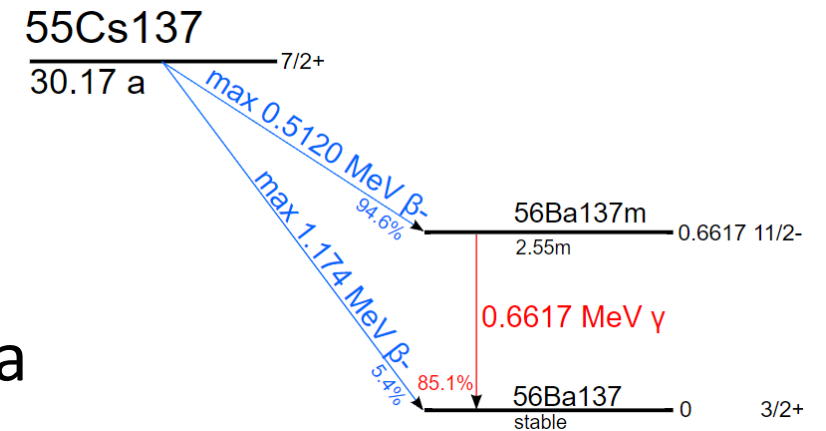
- Compton region:
Electrons recoil at different angles
- Compton edge:
Electron recoiled at 180° degree, maximum energy transferred to electron and deposited in the scintillator
- Multiple Compton scattering
- Photopeak: photon deposits all energy by photoelectric mechanism

Real spectrum

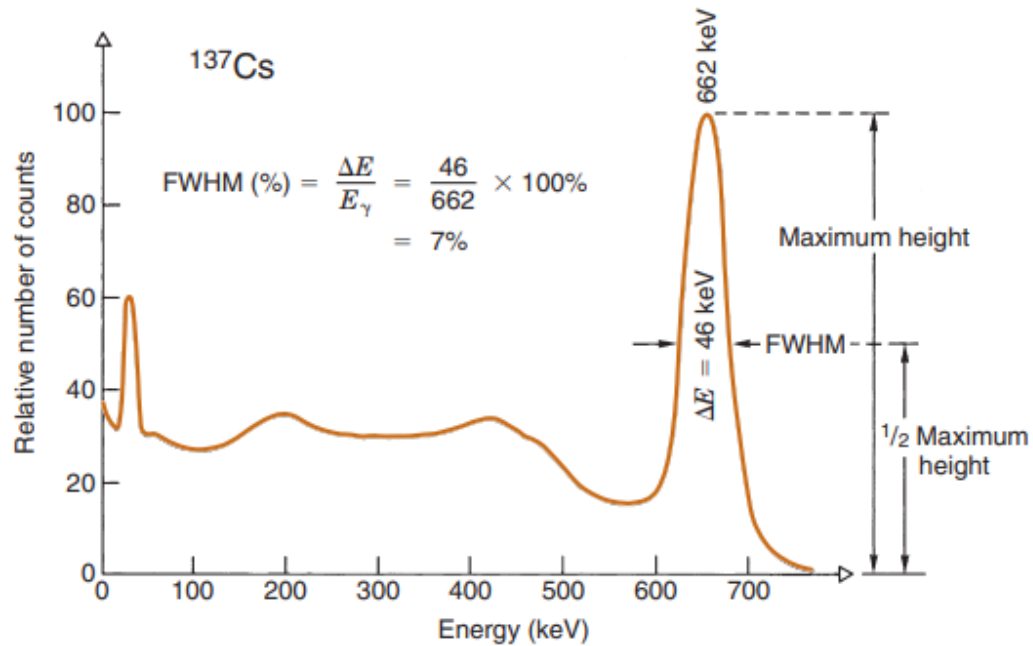


^{137}Cs decay emitting 662 keV gamma rays:

- Photopeak
- X-rays of Ba
- Backscattered peak



Spectrometry with NaI(Tl)



- Gaussian shape peak, instead of narrow-line.
- The width of the photopeak, ΔE , measured across its points of half-maximum amplitude is the energy resolution. This is referred to as the full width at half maximum.

$$\text{FWHM(\%)} = \frac{\Delta E}{E_\gamma} * 100\%$$

Detector efficiency

Efficiency with which detectors converts emission of a source into signal

$$\xi \left(\frac{\gamma \text{ rays}}{\text{sec}} \right) = A \left(\frac{\text{dis}}{\text{sec}} \right) * \eta \left(\frac{\gamma \text{ rays}}{\text{dis}} \right)$$

$$R = D \xi$$

Parameters:

- ξ rate of emission
- A activity
- η gamma per disintegration
- R rate of counts
- D efficiency

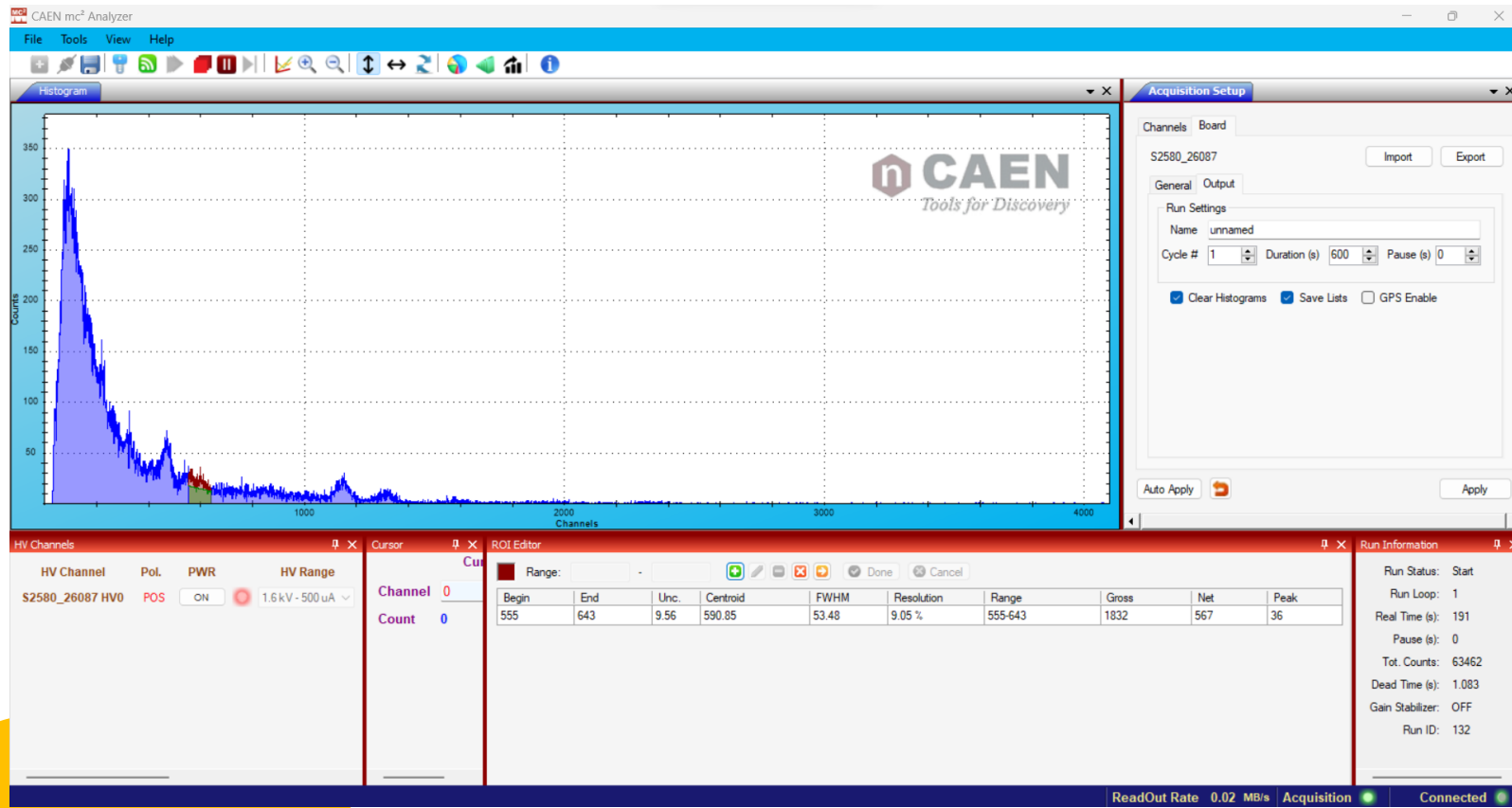
Multi-channel analyzer

Energy selection is made by multichannel analyser, allowing the entire spectrum to be analysed

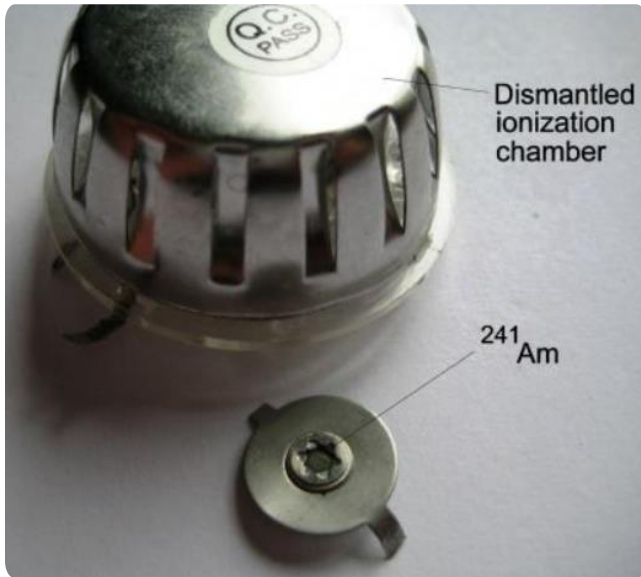
- Multi Channel Analyzer:
 1. Receives in input PMT output signals
 2. ADC digitizes the signals
 3. Produces the pulse-height histogram = plot of number of events from the PMTs as function of output amplitude

Channel = specific energy range → number of channels can be >1000
→ complete energy spectra produced

Gamma laboratory with MCA



1. Calibrate MCA with sources
2. Measure activity
3. Attenuation coefficient measure



Source

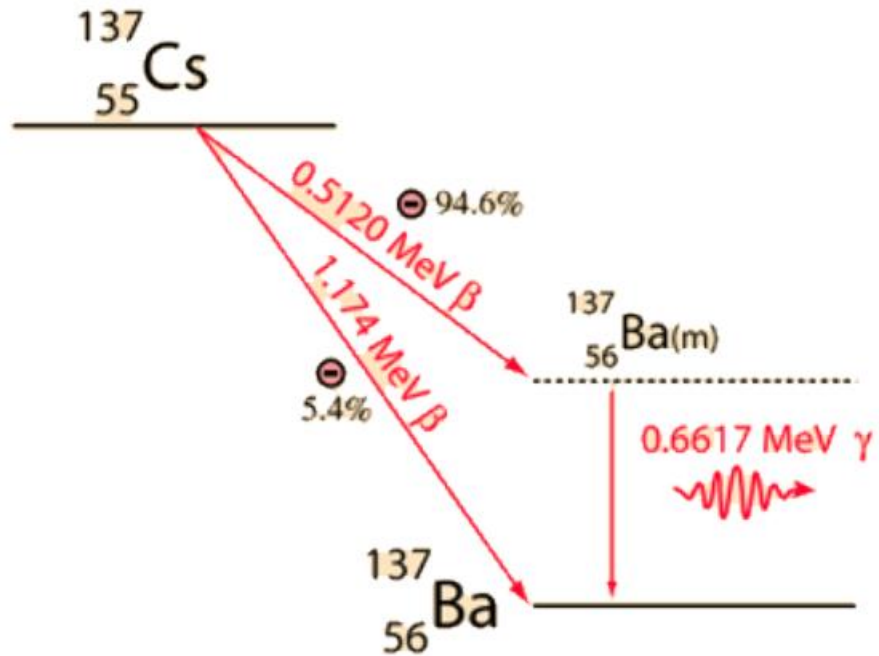
^{241}Am

Americium-241 is the most important radioisotope of americium from the point of view of the occurrence in environment. The other long-lived isotope ^{243}Am is produced in nuclear reactors in smaller activity compared to ^{241}Am . The activity of $^{242\text{m}}\text{Am}$ (half-life 160 years) that originated in nuclear weapons tests was nearly six orders of magnitude lower in comparison with ^{241}Pu activity from which ^{241}Am in-grows. Americium-241 is produced in nuclear power plants during activation of ^{239}Pu and ^{240}Pu by neutrons, which is followed by beta decay of ^{241}Pu ($T_{1/2} = 14.35$ years).

Americium -241 is used in many smoke detectors for homes and business, for measure levels of toxic lead in dried paint samples, to ensure uniform thickness in rolling processes like steel and paper production, and to help determine where oil wells should be drilled.

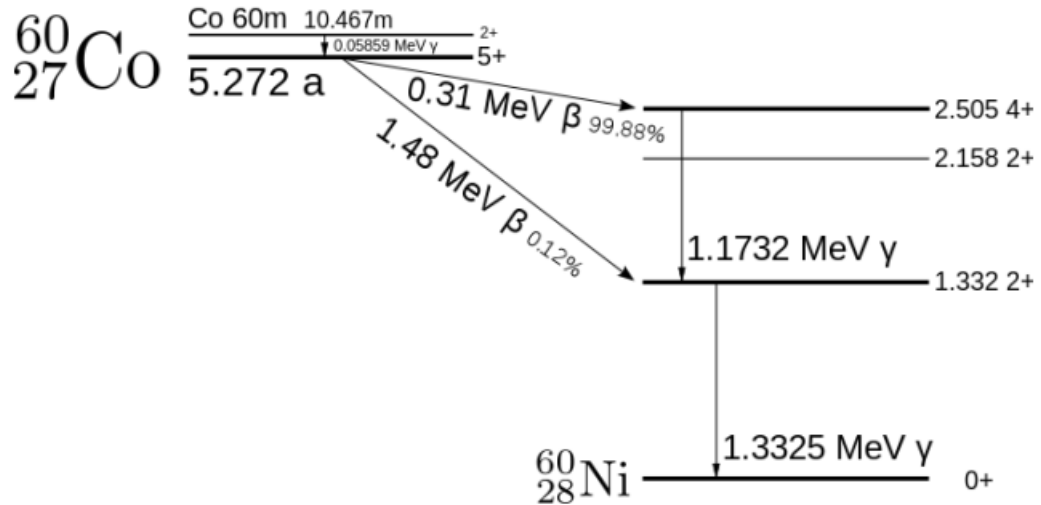
Source

^{137}Cs



Cesium-137 (Cs-137), or radium cesium is a radioactive isotope of cesium which is formed as one of the most common fission products from fission nuclear uranium-235 and other fissile isotopes in nuclear reactors and weapons nuclear. It is among the most problematic of the fission products with a medium-short half-life because it spreads easily in nature, due to the high water solubility of the most common chemical compounds of cesium. Small quantities of cesium-134 and cesium-137 were released into the environment during nearly all nuclear weapons tests and some nuclear accidents. Cesium 137 has a half-life of approximately 30.17 years. About 95 percent decays by beta emission to a nuclear isomer barium metastable: barium-137m (Ba-137m). The rest populate directly the ground state of barium-137, which is stable. Ba-137m has a half-life of approximately 153 seconds and is responsible for all emissions of gamma rays in cesium-137 samples. One gram of cesium-137 has a activity of 3,215 terabecquerels (TBq). The photon energy of Ba-137m is 662 keV

Source



^{60}Co

Cobalt-60 is a synthetic radioactive isotope of the metal cobalt. Due to its short average life of 5.27 years, cobalt-60 is not found in nature. It is produced artificially by activation neutronics of cobalt-59. Cobalt-60 decays by beta decay negative in the stable isotope nickel-60. The energized nickel-60 core emits two beams gamma with energies of 1.17 and 1.33 MeV per become nickel-60 stable

Energy (MeV)	μ/ρ (cm ² /g)	μ_{en}/ρ (cm ² /g)
1.00000E-03	2.911E+03	2.905E+03
1.50000E-03	9.536E+02	9.513E+02
2.00000E-03	4.206E+02	4.192E+02
3.00000E-03	1.288E+02	1.279E+02
4.00000E-03	5.466E+01	5.398E+01
5.00000E-03	2.792E+01	2.737E+01
6.00000E-03	1.608E+01	1.561E+01
8.00000E-03	6.750E+00	6.370E+00
1.00000E-02	3.481E+00	3.153E+00
1.50000E-02	1.132E+00	8.668E-01
2.00000E-02	5.798E-01	3.462E-01
3.00000E-02	3.009E-01	9.972E-02
4.00000E-02	2.304E-01	4.695E-02
5.00000E-02	2.020E-01	3.082E-02
6.00000E-02	1.868E-01	2.508E-02
8.00000E-02	1.695E-01	2.247E-02
1.00000E-01	1.586E-01	2.297E-02
1.50000E-01	1.406E-01	2.567E-02
2.00000E-01	1.282E-01	2.772E-02
3.00000E-01	1.111E-01	2.990E-02
4.00000E-01	9.947E-02	3.073E-02
5.00000E-01	9.079E-02	3.093E-02
6.00000E-01	8.395E-02	3.079E-02
8.00000E-01	7.372E-02	3.005E-02
1.00000E+00	6.628E-02	2.909E-02
1.25000E+00	5.927E-02	2.780E-02
1.50000E+00	5.395E-02	2.657E-02
2.00000E+00	4.630E-02	2.444E-02
3.00000E+00	3.715E-02	2.135E-02

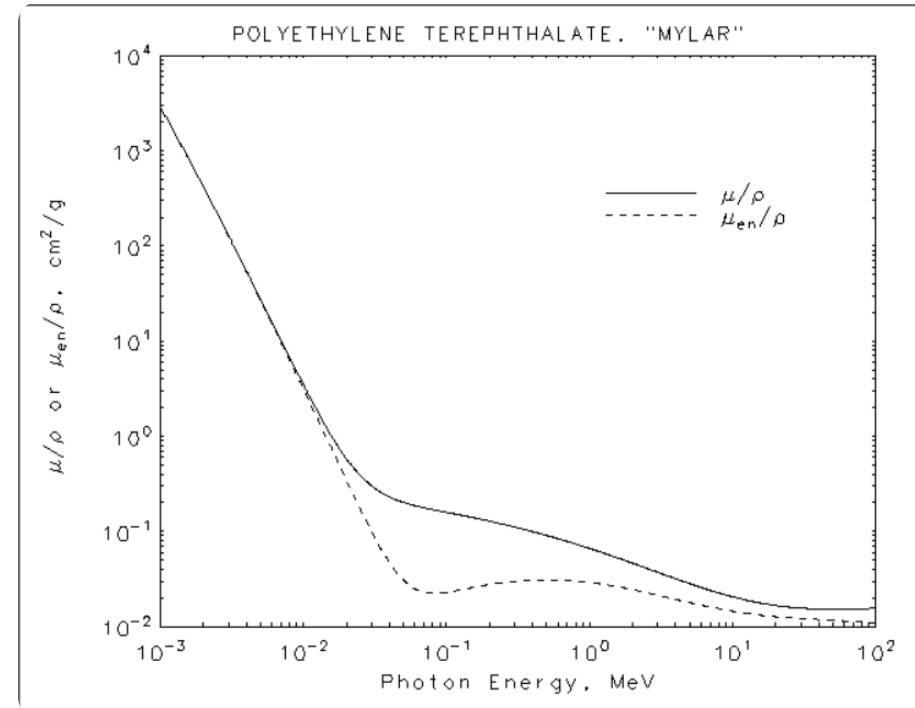
Americium →

Cesium →

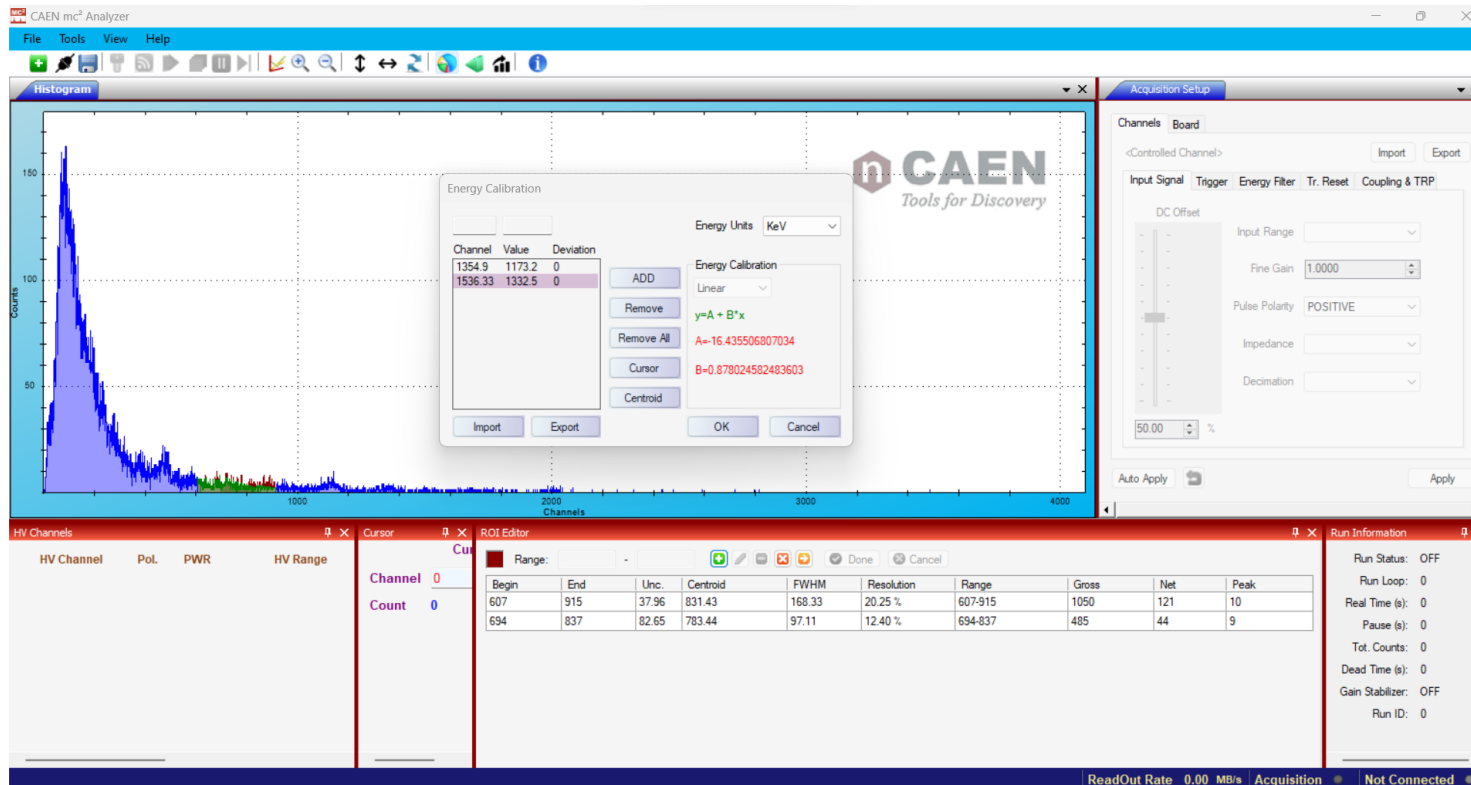
Cobalt →

NIST database

- Table shows mass attenuation coefficient at different energy for the polyethylene terephthalate

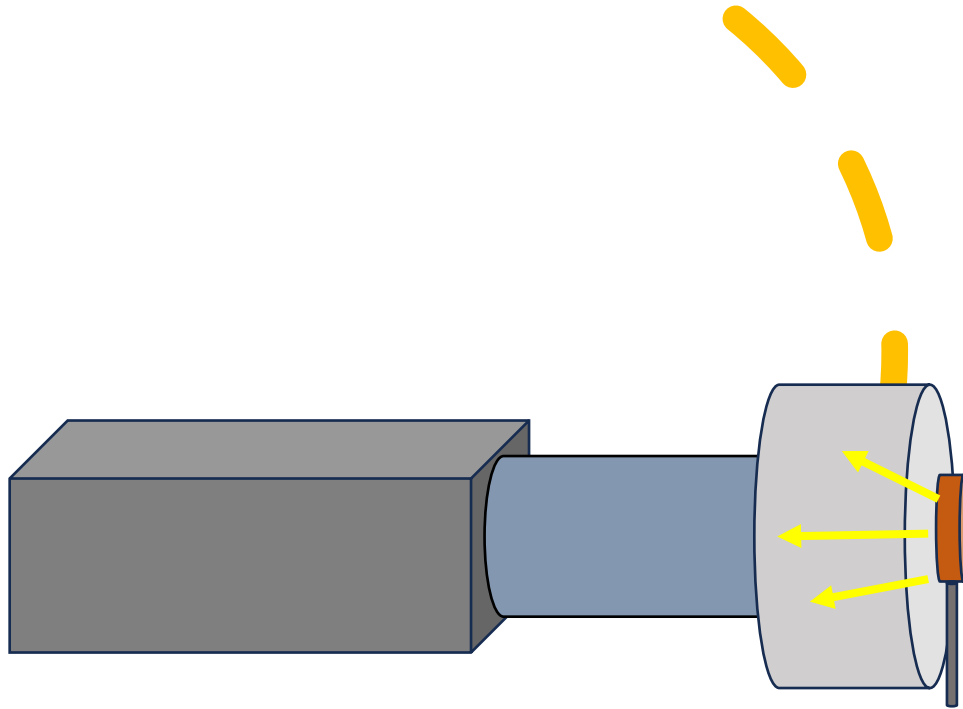


Calibration setup



Open calibration setup and set ^{60}Co peaks channels

Activity measure



$$R = D A$$

$$N = D A \Delta t$$


$$\frac{N}{D \Delta t} = A$$

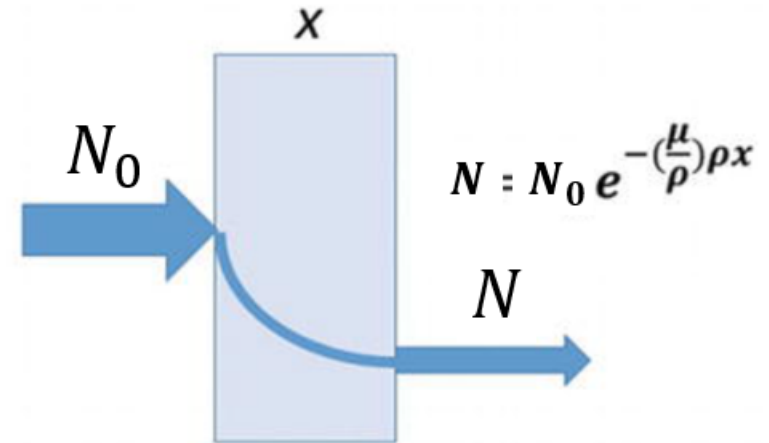
If we assume D is equal to 50% because of geometry, obtain A



Theory

Lambert-Beer Law

Source 



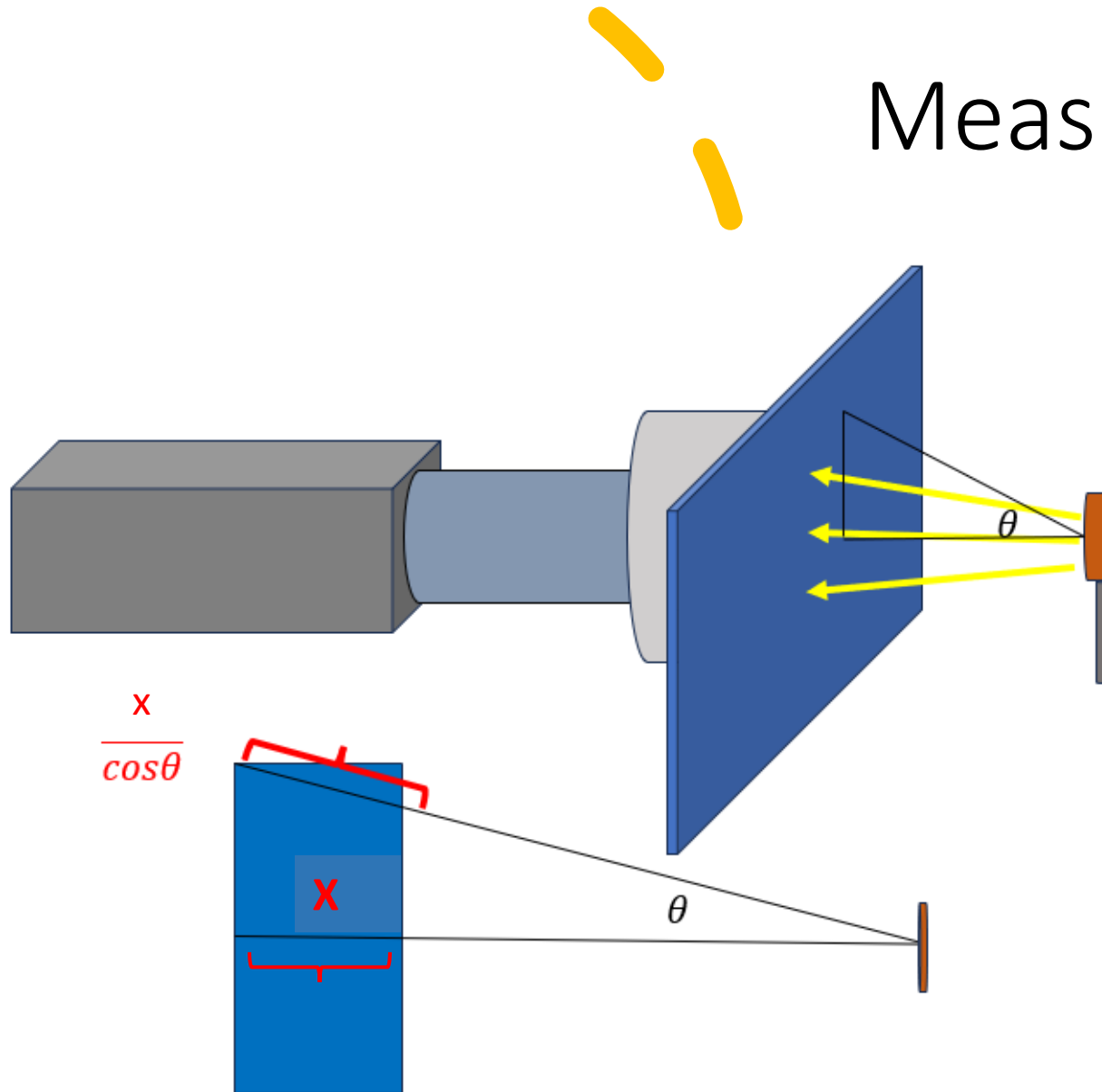
$$N = N_0 e^{-\mu x}$$

$N = \# \text{gamma transmitted}$

$N_0 = \# \text{gamma from source}$

$x = \text{thickness}$ $\mu = \text{attenuation coefficient}$

Measurements



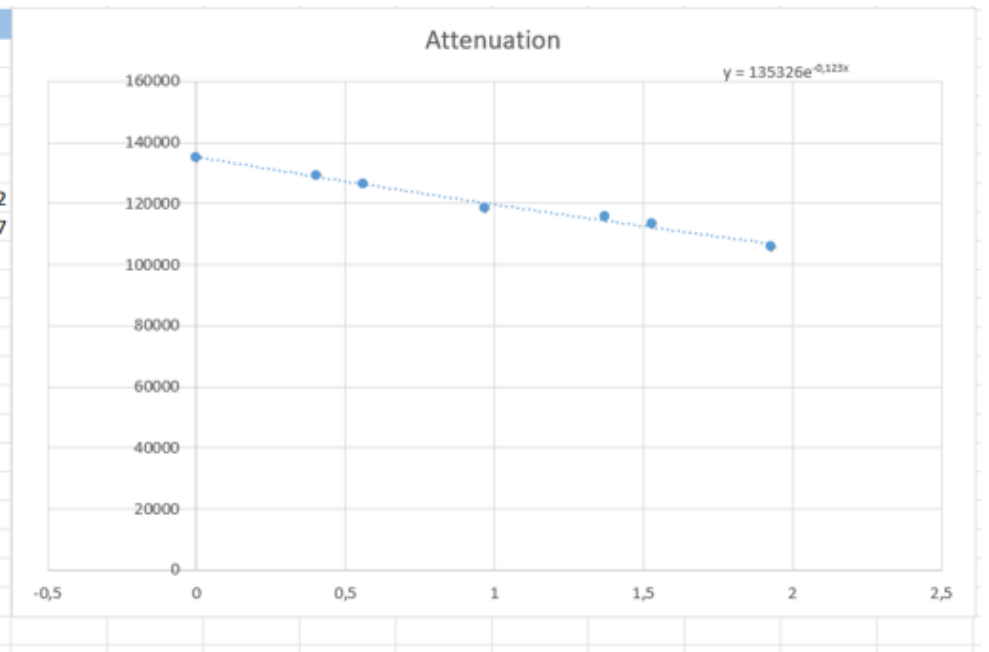
Measure the distance from the source to the detector, measure the thickness of the slice, position the slice in front of the detector

Measure the angle between the the source and the edge of the detector, like in figure.

$$\langle X \rangle = \frac{1}{2} \left(1 + \frac{1}{\cos\theta} \right)$$

Analysis

MEASURE OF ATTENUATION COEFFICIENT						
Shield	I	x (cm)	x corrected (cm)	sigma I	sigma x (cm)	distance (cm)
no shield	134978	0	0	367,3935	0,001	10
shield1	129355	0,387	0,402196473	359,6596	0,001	
shield2	126537	0,539	0,560165113	355,7204	0,001	theta 22
shield3	118536	0,931	0,967557923	344,2906	0,001	correction 1,039267
shield1+shield3	115643	1,318	1,369754395	340,0632	0,001	
shield2+shield3	113298	1,47	1,527723036	336,5977	0,001	
shield1+shield2+shield3	105802	1,857	1,929919509	325,2722	0,001	
polyethylene therephtalate (mylar)						
	0,1158					
polymethyl methacrylate						
	0,1035					
polyvynil chloride						
	0,1166					



Try with and without correction factor equal $\langle X \rangle = \frac{1}{2} \left(1 + \frac{1}{\cos\theta} \right)$