III. Energy Deposition in the Detector and Spectrum Formation

a) charged particles

Bethe-Bloch formula

$$-\frac{dE}{dx} = \frac{4\pi q_e^4}{m_0} \frac{z^2}{v^2} \cdot NZ \cdot \left[\ln \frac{2m_0 v^2}{I} - \ln(1 - \beta^2) - \beta^2 \right]$$

z, v: atomic number and velocity of projectile

N, Z: particle density and atomic number of absorber



⁽from *Particle Data Book*)

Helmuth Spieler

At $\beta\gamma \approx 3$ the differential energy loss assumes a minimum (minimum ionizing particles), independent of absorber.



dE/dx vs. *E* of protons in silicon

minimum ionization at 2 - 3 GeV





Minimum ionization at ~1 MeV.

At electron energies >50 MeV radiative energy loss dominates, so the stopping power increases more rapidly beyond the energy of minimum ionization than for protons. For small velocities ($\beta <<1$)

$$-\frac{dE}{dx} \propto \left(\frac{z}{v}\right)^2 \cdot NZ$$

 $\Rightarrow dE/dx$ increases with projectile charge squared decreases with increasing projectile energy (v^2)

increases with target atomic number and density

dE/dx for alpha particles in various materials



(from Knoll)



dE/dx vs. *E* for various charged particles in Si

 $- \frac{dE}{dx} (\text{keV}/\mu\text{m})$

50

10

5

1

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2



(from Knoll)

Alphas

³He⁻

Tritons

Deuterons

20

Protons

10

5

Particle energy (MeV)





(from Northcliffe and Schilling, Nucl. Data Tables A7 (1970) 233)

Note that the vertical scale is normalized to Z^2 and density ρ .

In contrast to particles of low atomic number, the specific energy loss of heavy particles initially increases with energy. With increasing velocity, orbital electrons are stripped from the nucleus, thereby increasing the effective charge of the projectile. Beyond the peak dE/dx, where the projectiles are fully stripped, the specific ionization falls as predicted by the Bethe-Bloch formula and attains minimum ionization at 1 GeV/amu.

Charged particles have a well-defined range (stopping distance)



Range-Energy curves for various charged particles in Si

(from Knoll)

For electrons, however, the range of individual particles is subject to substantial fluctuations due to scattering at low energies (straggling).



RANGE OF ELECTRONS IN SILICON

For electrons the range expressed in g cm⁻² is practically the same in all absorbers.

$$r[g \operatorname{cm}^{-2}] = r[\operatorname{cm}] \cdot \rho[g \operatorname{cm}^{-3}]$$

If the range-energy curve is known for one material, it can be translated to other materials by simple scaling.

$$r_1 \rho_1 = r_2 \rho_2$$

Towards the end of their range the specific energy loss of charged particles increases and they move more slowly, depositing even more energy per unit distance.



For a beam of particles the Bragg peak is smeared because of multiple scattering.

At low energies the range of particles decreases drastically with increasing projectile charge.

For $E=$ 5 MeV in Si:	р	$R=$ 220 μ m
	α	<i>R</i> = 25 μm
	¹⁶ O	<i>R</i> = 4.3 μm
	⁴⁰ Ca	<i>R</i> = 3.0 μm
	¹³² Xe	<i>R</i> = 2.0 μm
	¹⁹⁷ Au	<i>R</i> = 1.4 μm

⇒ "Bragg Peak"

Energy Loss of Neutrons

Cross section much smaller (no Coulomb interaction).

Energy deposition due to knock-on collisions that displace absorber atoms, which deposit energy as charged particles.

Maximum recoil energy transferred to a nucleus of mass number A for head-on collision

$$E_{\max} = \frac{4A}{\left(A+1\right)^2} E_n$$

Maximum for protons ($E_{max} = E_n$), whereas for C $E_{max} = 0.35 E_n$

Hydrogenous absorber most effective as neutron detectors

 \Rightarrow plastic scintillators

For fast neutrons up to 10 MeV the scattering is isotropic in the center-of-mass system, so the energy distribution of the recoil protons is uniform from $E_p = 0$ to $E_p = E_n$.



Except for very low energy photons, the spectrum provided by plastic scintillators is dominated by Compton scattering.

Interactions of Gamma Rays

In contrast to charged particles, which deposit energy continuously along their track, photon interactions are localized.

In passing through a medium, photons will traverse a certain distance unaffected, until depositing energy either by

- a) photoelectric absorption
- b) Compton scattering
- c) pair production

The probability of undergoing an interaction is an exponential function of distance. The fraction of photons that suffered any interaction after traversing a distance x is

$$f = 1 - \exp(-\mu x)$$

where μ is a linear absorption coefficient, expressed in cm⁻¹, which is the sum of the individual absorption coefficients of the relevant interactions.

The absorption can be parameterized more generally by the mass attenuation coefficient μ / ρ , expressed in cm²g⁻¹, which is independent of the density or physical state of the absorber.

For a mixture of elements of weight fractions w_i the composite mass attenuation coefficient is

$$\frac{\mu}{\rho} = \frac{\mu_1}{\rho_1} w_1 + \frac{\mu_2}{\rho_2} w_2 + \dots$$

a) Photoelectric Absorption

Photon deposits its total energy in a single interaction with an absorber atom.

A photoelectron is emitted with the energy

$$E_e = E_{\gamma} - E_b$$

where E_b is the binding energy of the photoelectron.

Photoelectric absorption dominates at rather low energies, but the cross section increases rapidly with Z.

$$\sigma_{pe} \propto \frac{Z^n}{E_{\gamma}^{3.5}}$$

where n varies from 4 to 5.

At energies and absorber atomic numbers where this process is important, the photoelectron typically is absorbed within a short distance, so the total energy is registered in the detector.

X-rays emitted by the decay of the ionized atom to its ground state are also absorbed in the detector.

 \Rightarrow measured spectrum



(from Knoll)

b) Compton Scattering

Photon scatters off of an electron.

 \Rightarrow photon deflected with decreased energy

$$E_{\gamma} := \frac{E_{\gamma}}{1 + \alpha (1 - \cos \Theta)}$$

where

$$\alpha \equiv E_{\gamma} / m_0 c^2$$

Relative number of photons Compton scattered per unit solid angle vs. scattering angle Θ and initial photon energy.



(from Knoll)

The primary photon is incident from the left.

At higher energies photons scattered near the far boundary of the detector will tend to escape from the detector, leading to a deficit in the measured energy.



$$T_{cm} = \frac{E_{\gamma}}{1 + (1/2\alpha)} ,$$

(from Birks)

If the scattered photon exits the sensitive volume of the detector without interaction, the electron energy spectrum is what will be registered by the detector. Electron scattering at $\Theta=\pi$ imposes an upper bound on the energy of the scattered photon

$$E_C = E_{\gamma} - E_{el}|_{\Theta = \pi} = \frac{E_{\gamma}}{1 + 2E_{\gamma} / m_0 c^2} ,$$

the "Compton edge".



(from Knoll)

For large photon energies $E_{\rm y}$ this energy converges toward a constant value, placing the Compton edge below the primary photon peak by

$$E_C(E_\gamma \rightarrow \infty) = \frac{m_0 c^2}{2} = 256 \text{ keV}$$

c) Pair Production

If the photon energy is twice the mass of the electron, electronpositron pairs can be produced.

No simple expression for the cross section vs. energy, but this process becomes significant at photon energies above several MeV.

The excess photon energy beyond $2m_0c^2$ goes into kinetic energy shared between the electron and positron.

The positron can interact within the detector yielding two 511 keV annihilation photons

If these are absorbed in the detector, the measured energy spectrum is



(from Knoll)

Absorption Processes vs. Energy in NaI



Relative importance of the three major type of gamma-ray interactions vs. energy and atomic number Z of the absorber.



The lines show the values of *Z* and $E_{\gamma} = hv$ where the two neighboring effects are equal.