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## **GEANT4 Radioprotection studies of the ERHA system for protontherapy treatment**

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## **Abstract**

*This work is focused on the radiation protection for a protontherapy facility. The aim is to simulate with the best accuracy the prompt radiation field of the proton accelerator situated in Ruvo di Puglia, owned by Linearbeam s.r.l. company. In order to simulate it, is used Geant4, a software for interaction simulations of particles with matter. Thanks to internship work, thesis speaks about cancer therapy with a new method for particle acceleration, a linear beam. For a complete overview of the therapy, this work starts with a crash course on interactions of particle with matter, goes specifically to biological matter, then is shown a brief introduction to shielding studies for a particle acceleration facility, and then a presentation of Geant4. At the end, the main aspects of the proton accelerator are simulated, from proton hitting material of beam-pipe to detectors used to measure dose.*



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# Introduction

Cancer is a large group of diseases that can start in almost any organ or tissue of the body when abnormal cells grow uncontrollably, go beyond their usual boundaries to invade adjoining parts of the body.

Cancer is the second leading cause of death globally, accounting for an estimated 9.6 million deaths, in 2018 [1]. There are numerous methods of treatment, like for examples surgery, chemotherapy, immunotherapy, radiotherapy ( $\approx 50\%$  of all cancer patients) and hadrontherapy, also called Ion Beam Therapy.

Hadrontherapy had a rapid development in recent years due to the progresses in technology and radiation oncology techniques. In the last 50 years there was a continuous expansion, at the end of 2016 there were 61 proton centres and 9 heavy ion centres, with 149345 patients treated with proton and 21580 patients treated with  $^{12}\text{C}$ .

Hadrontherapy represents a valid alternative to the conventional radiotherapy that uses photons or, more rarely, electrons. Comparing these therapies, hadrontherapy advantages comes from the different lost energy mechanism. The dose release profile of radiation with photons presents a peak at short distance from the patients's skin followed by a decreasing release of the radiation in accordance with the absorption law. On the other hand, hadrontherapy presents a low dose profile at the beginning of the path and a sharp maximum, *Bragg peak*, near the end. This therapy provides a high irradiation accuracy of the tumor volume and a reduced damage to the surrounding healthy tissues. The particles involved in this therapy present high capability of inducing a direct damage to the DNA of cancerous cells.

Unfortunately, when using these beams, there is a major increase in the presence of fragments derived from the nuclear interactions of the beam and the patient tissues. Consequently a dose is released in the entry region and beyond the Bragg peak, still not completely studied. Proton treatments are the most widespread but even in their employment occurs the problem of fragmentation.

FOOT (FragmentatiOn Of Target) experiment was designed with purpose of providing for the lack of experimental measurements of nuclear reaction cross sections for fragments produced in the interaction between tissue nuclei and charged particles of the beam, with the task of study both projectile and target fragmentation where the latter was neglected by previous experiments. Fragmentation problem vanishes for protontherapy, because proton interaction are mostly due to ionization.

In Chapter 1 is described how particles interact with matter, with an overview of the

most probable mechanisms. In Chapter 2 is presented an introduction to the effects at different energies, for different particles, of the interaction of particles with biological matter, focused on what happens when radiation is used to kill cancers.

In Chapter 3 is described the Linearbeam accelerator facility, an overview of the ERHA project, a detailed description of the components. In Chapter 4 is explained how to shield from steady-state radiation fields, with a focus on the protection study of an accelerator facility, involving some fundamentals concepts and quantities. In Chapter 5 is shown the GEANT4 simulation work, main object of this thesis, in which in three steps is built a very similar model of the accelerator facility in order to obtain the gamma and neutron dose distribution inside bunker.

In Chapter 6 are reported measured quantities, dosimeter and set-up description, in order to compare them with simulation.

# Chapter 1

## Radiation interaction with matter

A brief overview of the physical process that occurs when charged and neutral particles cross matter is given in this chapter. Charged particles interact via Coulomb force and nuclear fragmentation, while neutrons could scatter with nuclei and produce secondary charged particles.

### I. CROSS SECTION

The cross section is a measurement of the probability that a reaction happens. Let's imagine a diffusion experiment in which a beam of particles smashes on the surface of the target material and interacts all along the thickness  $\delta x$ , like in figure 1.1[2].  $N_i$  is the number of particles of the beam, and  $n_b$  is the density of scattering centers, so the product  $n_b \delta x$  is the number of scattering centers for a unitary surface. Now supposing that each particle in the beam can interact on average with at most one and only one particle of the target, the number of reactions in unit of time  $\frac{dN_r}{dt}$  is proportional to number of particle of the beam in the same unit of time, and  $n_b \delta x$ . In order to write the proportionality:

$$\frac{dN_r}{dt} = \sigma \frac{dN_i}{dt} n_b \delta x \quad (1.1)$$

So  $\sigma$  is like a surface, and it is measured in *barn*, 1 *barn* corresponds to  $10^{-24} \text{cm}^2$ .

### II. INTERACTION OF CHARGED PARTICLES

Particles with electric charge could interact with electrons and nuclei, depending on cross section: when a particle interacts with the entire atom cross section is much bigger than when it interacts with the nucleus, so the most probable process happens with atoms, and for the sake of this thesis it is possible to distinguish between three main phenomena: *bremstrahlung*, cherenkov and ionization.



i. Bremsstrahlung

Bremsstrahlung consists of light emission from a charged particle of mass  $m$  that interacts with an electromagnetic field, specifically the  $Z$  charged nucleus's field, equal to:

$$-\frac{dE}{dx} = \frac{4N_a Z^2 \alpha^3 (hc)^2}{m^2 c^4} E \ln\left(\frac{183}{Z^{1/3}}\right) \sim \frac{Z^2}{m^2} \quad (1.2)$$

This contribution is negligible for protons and ions due to  $\frac{1}{m^2}$ .

ii. Cherenkov

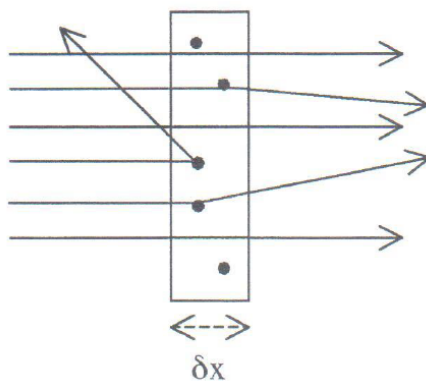
This phenomena is due to light emission that occurs if a charged particle travels a medium faster than speed of light in that medium. The cause of this emission is linked to the polarization and depolarization of matter when particle crosses it. Every point of the trajectory gives rise to a spherical wave front, with  $v = c/n$ . So the threshold for which phenomena happens is  $\beta > 1/n$ .

iii. Ionization

Charged particles main process is the ionization or excitation of atoms, and the mean rate of energy lost per unit length of material is called *stopping power* and it is expressed by the Bethe-Block formula:

$$-\left\langle \frac{dE}{dx} \right\rangle = \frac{2\pi n_e r_e^2 m_e c^2 z^2}{\beta^2} \left[ \ln\left(\frac{2m_e c^2 \beta^2 T_{max}}{I^2(1-\beta^2)}\right) - 2\beta^2 + 2zL_1(\beta) + 2z^2L_2(\beta) - 2\frac{C}{Z} - \delta + G \right] \quad (1.3)$$

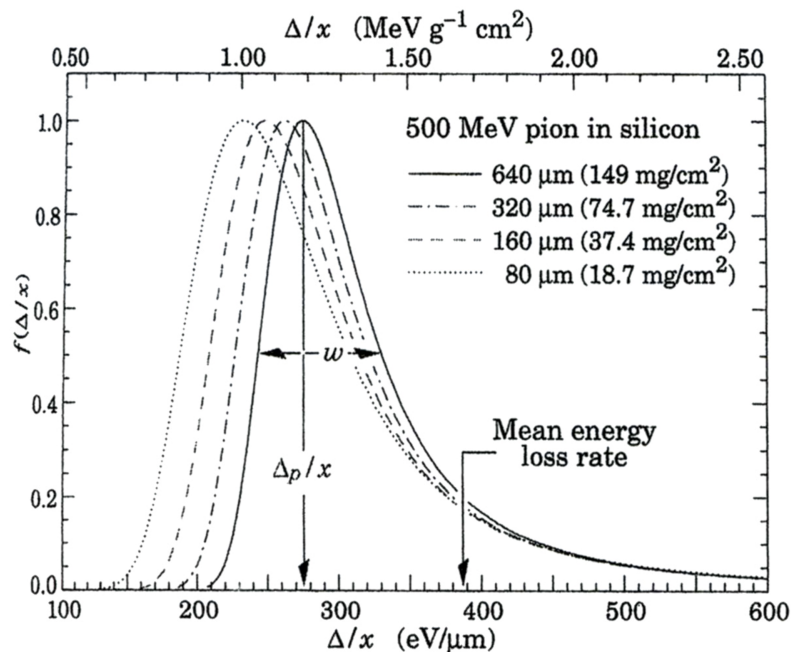
- $\beta$  charged particle velocity in c units
- $I$  mean excitation energy, material-dependent



**Figure 1.1:** Diffusion experiment on a target.

- $\delta$  density correction
- $C$  is the shell correction, important at low energies
- $T_{max}$  maximum energy transfer to an electron (from kinematics)
- $L_1$  Barkas correction ( $z^3$ ) responsible for the difference in stopping power for particles-antiparticles
- $L_2$  Bloch ( $z^4$ ) correction
- $G$  Mott corrections

The formula is valid for particles with mass bigger than the electron mass. The *Barkas effect* was discovered by Walter Barkas, who saw that positive and negative pions with same energy that went through emulsion material had different ranges. This difference relies on the sign of the electromagnetic force, positive particles bring near electrons while negative particles repel them, changing the mean charge density of the material. The energy lost shows a distribution that depends upon the thickness of the material. For thick materials a large number of ions was created (from 30000 to 70000 ions pair for  $\alpha$  particle in air), leading to a Gaussian distribution. For thin materials, the energy straggling in statistical fluctuations depends upon a little number of interactions with a huge loss of energy. This distribution is called Landau 1.2. In Figure 1.3 there is a plot of BetheBlock for different materials.



**Figure 1.2:** Plot of the Landau distribution, where  $\Delta$  is the energy loss and  $x$  is the thickness[3].

This shows also that particles lose their energy for very low energy values, so at the end of the *Range*. This quantity is the mean length of a particle inside materials. In order to obtain an analytic formula for the *Range* a method is integrate over all the deposited

energy, but it's not easy.

$$R = \int_0^{T_0} \left( -\frac{dE}{dx} \right)^{-1} dE \quad (1.4)$$

This is the formula for the range, where  $E_0$  is the starting energy. This approach neglects that energy losses are stochastic in nature and that secondary particles possess an energy and range distribution. In addition, electrons do not have a straight path but suffer multiple scattering with many changes in direction [4]. In order to simplify the integral, it could be useful to use the *Bragg-Kleeman approximation*, under the condition of a *Continuous Slowing Down Approximation* (CSDA) in which particles lose their energy slowly and continuously. The condition led to a simplification in a very common situation, in which particles go through a compound with different stopping power, giving this formula:

$$\frac{dE}{d\chi} = \sum_i^N W_i \left( \frac{dE}{d\chi} \right)_i \quad (1.5)$$

in which the  $\chi$  is the massive length,  $\chi = \rho x$ , and  $W_i$  is the fraction of material in the compound.

**Massive particles** Particles that have got a mass bigger than electrons are heavy or massive, like protons or  $\alpha$ . They lose energy inside materials by ionization. When particles lose energy, their velocity decreases and lose more and more energy, with an ionization density peaked at the end of range, so the deposited energy quantity reached a peak, called *Bragg peak* (see Figure 2.6), where  $\beta \sim \frac{z^{2/3} v_0}{c}$ , and  $v_0 = \frac{e^2}{\hbar}$ . At peak is reached the maximum effect, as is explained in Chapter 2.

**Electrons and positrons** Electrons lose energy by Bethe-Block with a change, due to the identity between projectile and target particles. Bremsstrahlung overcomes ionization at a critical energy value, that depends on material, but in general it is comparable with ionization at  $10\text{MeV}$ .

### III. INTERACTION OF NEUTRAL PARTICLES

Neutral particles don't lose energy ionizing matter, processes are more complicated. First of all, it's fundamental to distinguish between photons and neutrons: photons haven't mass, so main interactions with ordinary matter is pure electromagnetic. Neutrons are massive particles that could interact with nuclei, by elastic or anelastic collisions.

#### i. Photons

Photons interact with matter at atomic and nuclear level, with different processes. First photon could be absorbed, giving rise to an electron, or it could scatter on an electron, while at nuclear level, a photon could disappear in a couple of electron-positron. These

particles go deeper into matter more than charged particles, and they don't lose energy smoothly but all the energy is lost. So, in order to understand a photon beam interaction, it is useful to derive the intensity of the beam:

$$I(x) = I_0 e^{-\mu x} \quad (1.6)$$

where  $I_0$  is the incident intensity,  $x$  is the thickness of material,  $\mu$  is the *attenuation coefficient*. So it is linked to the cross section:  $\mu = \frac{\rho N_A \sigma}{A}$ .

**Photoelectric effect** The photoeffect is the first demonstration of quantum physics, photon is absorbed by atom, and the electron ejected has got an energy:

$$E = h\nu - \phi \quad (1.7)$$

where  $\phi$  is the binding energy. The cross section over the threshold goes down as energy increases. Electron leaves a hole inside atom, that will be filled by an electron of another shell, and this process goes on in order to re-establish an equilibrium of electrons, giving rise to emission of photons. Cross section of photoeffect is  $\sigma \propto \frac{Z^{4.5}}{E^{3.5}}$

**Compton effect** When electrons have more energy and very small binding energy, they are free. A photon hitting these electrons may be scattered and lose energy, changing its frequency. The cross section is  $\sigma \propto \frac{Z}{E}$ .

**Pair production** Phenomena of pair production happens when energy of photons overcomes  $2m_e c^2$ . It's made possible by the presence of matter: a nucleus interaction that absorbs the momentum. Just a simplified description, but a very clear explanation is possible in the framework of Quantum Field Theory. Cross section is  $\frac{Z^2}{\ln E}$ . In order to visualize different interaction it is needed to define *mass attenuation coefficient*  $\frac{\mu}{\rho}$ , as showned in figure 1.4.

## ii. Neutrons

Neutrons could interact only by strong force with nuclei. This interaction is a short range force, so neutrons have to reach a very small distance from nuclei on order to interact, less than  $10^{-15}m$ . For this reason neutrons go very deep into matter. Neutrons should be divided by their energy, like in Table 1.1 All the possible interaction of neutrons with nuclei are listed below:

1. **Elastic Scattering**: main lost energy mechanism into MeV region.
2. **Anelastic Scattering**: nuclei is excited and decays with gamma or other type of decay. Always possible into MeV region.
3. **Radiative capture**:  $n + (Z, A) \rightarrow \gamma + (Z, A + 1)$ , and  $\sigma \propto \frac{1}{v}$ , with  $v$  velocity of neutron.

4. **Nuclear Reactions:** neutron is captured and other charged particles are emitted. Cross section goes as the radiative capture, but there could be some resonances, into a range from eV to keV.
5. **Fission:** this phenomena happens for thermal neutrons.
6. **Hadronic cascade:** over 300 MeV.

In many situation, like detectors or nuclear plants, a very important process to study is the *moderation* of neutrons, a method to decrease their energy. When they go inside matter, they scatter with nuclei before they reach thermal or cold energy. At this point it is very probable a capture reaction. A specific explanation of nuclear reaction in a detector is written in chapter 6.

#### IV. NUCLEAR FRAGMENTATION

The problem of hadrontherapy is that at energy of  $10^2$  MeV/nucleon, fragmentation is the most frequent nuclear interaction: the projectile collides with target ppheriferically, with few nucleons participating and products are separated in quasi projectile fragment and quasi target fragment both spectator of the interaction, while the region of interaction fragments in few nucleons like in Figure 1.5. The primary beam particle, like  $^{12}\text{C}$  nuclei, can produce lighter fragments which, with lower energy, will continue to move through the material releasing energy inside it. The dose released beyond the Bragg peak depends on the charge of the particle: it is smaller if particles are protons (around 15 %), larger if they are  $^{12}\text{C}$  nuclei and even more if  $^{20}\text{Ne}$  nuclei. Depending on impact parameter between the particle and other nuclei of the material we could have a situation with one single fragment or many much lighter fragments, depending on the impact parameter. Fragments from quasi-projectile have velocity almost equal to velocity of the beam and are emitted in a narrow angle and then those fragments have a larger range with respect to the beam, because they are lighter and have almost same velocity.

The other fragments from quasi-target have a wider angular distribution and lower energies so they will stop before the quasi-projectile. This is the case of light particles such as protons, deuterons and helium. The dose beyond the Bragg peak comes from the quasi-projectile contribution whereas the wide angular halo comes from the quasi-target

**Table 1.1:** Classification of neutrons by their energies.

Neutron	Energy
High Energy	$> 100$ MeV
Fast	$100$ keV $\div$ $100$ MeV
Epithermal	$100$ meV $\div$ $100$ keV
Thermal or Slow	$25$ meV
Cold or Ultracold	$1$ $\mu\text{eV}$ $\div$ $1$ meV

fragments which have larger lateral displacement.

It's possible to study with Monte Carlo simulation the informations about cross-sections and other relevant parameters. The goal is to study how many fragments are obtained and of which energy. An example of fragments distributions is in Figure 1.6. Fragments play an important role in spreading collateral damage to the healthy tissues crossed by the incident particles. Since the fragments have higher range and different directions, they cannot be neglected in treatment planning.

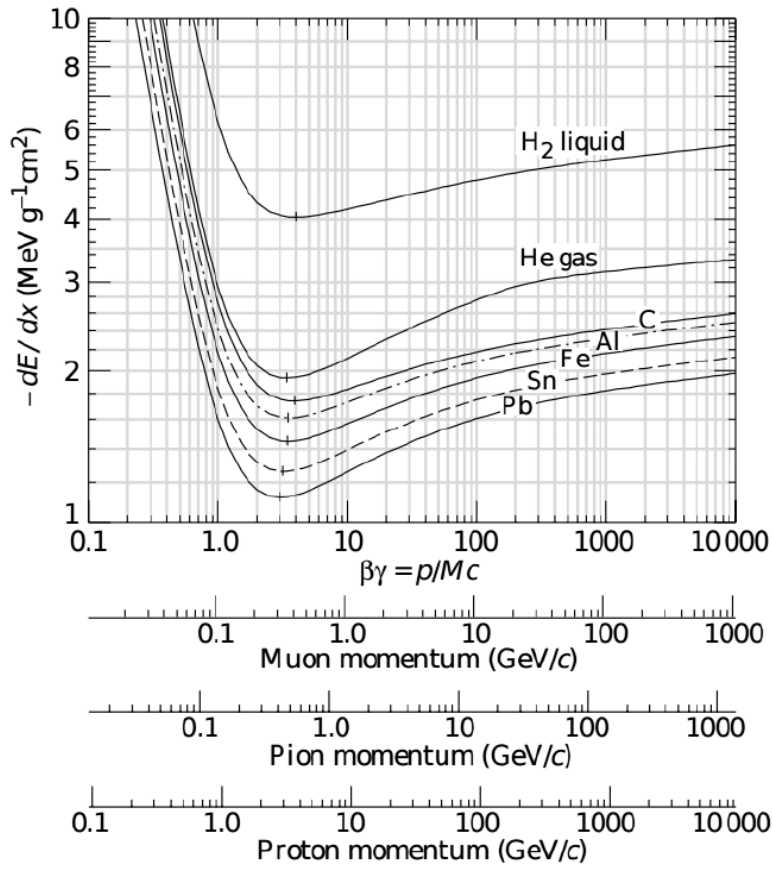


Figure 1.3: Plot of the BetheBlock formula[4].

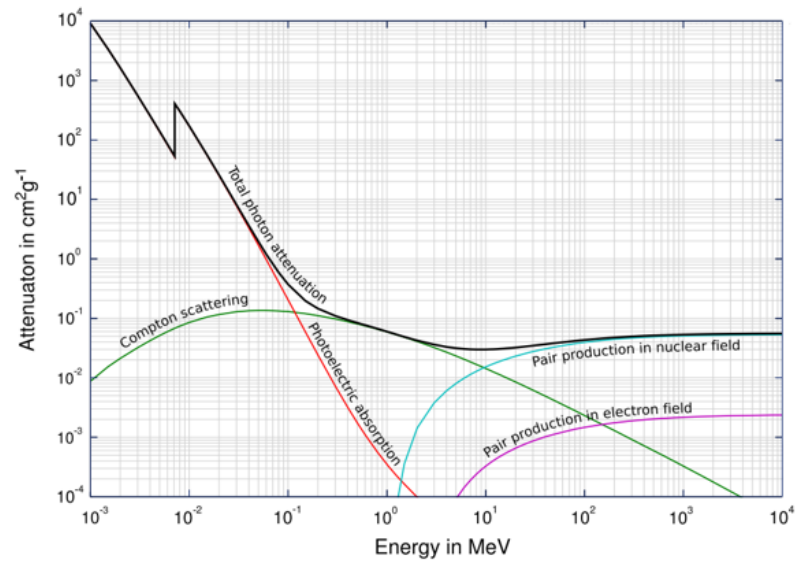


Figure 1.4: Plot of the attenuation coefficients for Fe [5].



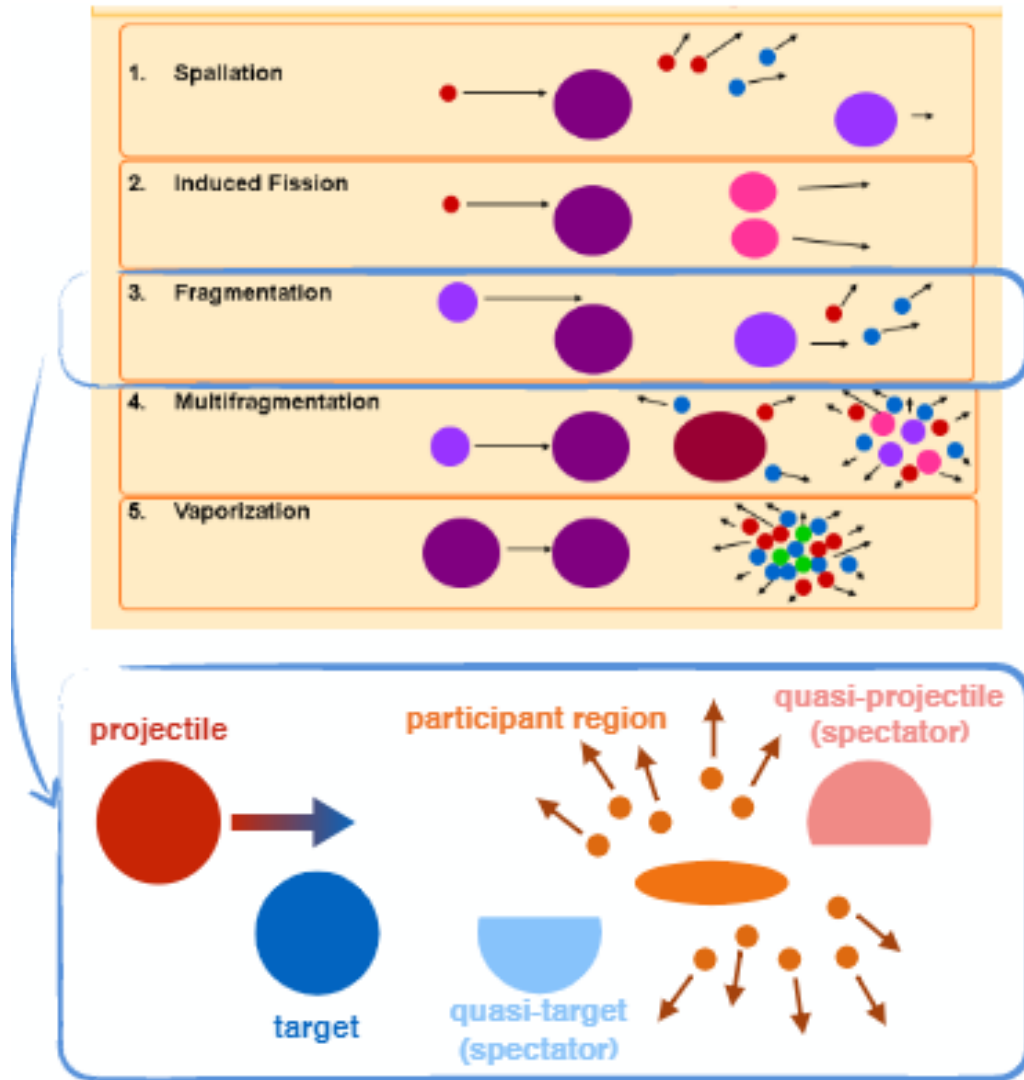
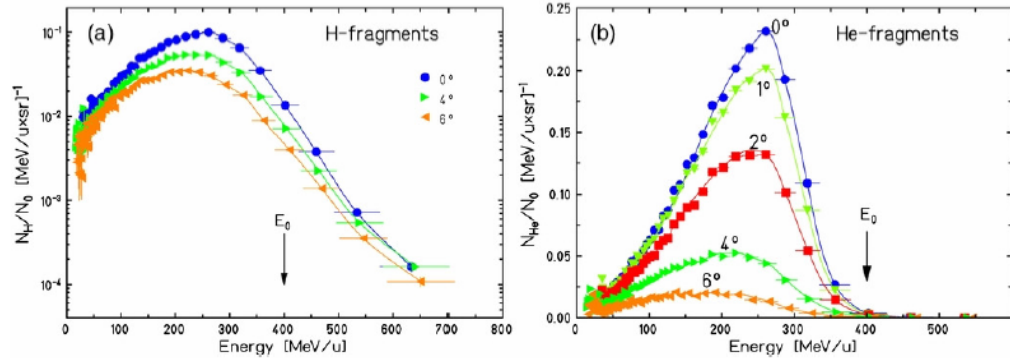


Figure 1.5: An illustration of the nuclear interactions with a zoom on the fragmentation.



**Figure 1.6:** Energy spectrum of secondary particles, Hydrogen (a) and Helium (b), produced in the fragmentation of a  $^{12}\text{C}$  beam at 400 MeV/u in a water absorber 27,9 cm deep; distinct curves correspond to different emission angles ( $0^\circ$ ,  $4^\circ$ ,  $6^\circ$  on the left and  $0^\circ$ ,  $1^\circ$ ,  $2^\circ$ ,  $4^\circ$ ,  $6^\circ$  on the right)[6].



## Chapter 2

# Radiation biology

Biological radiation actions comprises all levels of biological organization. The interaction of radiation and biological system is as old as life itself, it certainly played a central role in evolution of self-organizing structures. The first effect of radiation action on biological systems is the transfer of energy to essential cellular components. This requires the introduction of different concepts and quantities.

### I. BIOLOGICAL RADIATION EFFECTS

Ionizing radiations are made of particles that ionize atoms while they cross matter. The interaction of these particles, charged or not, light or heavy depends upon several factors, like energy. Photons with MeV energy can mainly do Compton and pair production and produce electrons and positrons of not so high energies which are light particles so they would lose energy through Bethe-Block. On the other hand a MeV neutron can interact with nuclei scattering on them. Nuclei are charged more than one so they will lose their energy in a small portion of tissue.

#### i. Dose

Every physical or biological effect induced in the tissue is a consequence of the energy deposited in the matter by radiation. It's important to define the quantity of energy deposited by radiation per unit mass: the absorbed dose

$$D = \frac{dE}{dm} \quad (2.1)$$

defined as the expectation value of the absorbed energy divided by the mass of the volume. The unit of dose is the *Gray* (Gy) which equals 1 J/kg. An older and officially outdated unit is the *rad* (rd):

$$1rd = 100erg/g = 0.01Gy \quad (2.2)$$

**Equivalent dose** The radiation received corresponds to dose, but different exposure (electrons, protons, neutrons, gamma, ...) results into different physical damages. Stochastic effects occurrence depends not only on the amount of energy absorbed, but also on the nature of the radiation generating the dose. This difference is taken into account by weighting the absorbed dose by a factor related to the quality of the radiation, called the radiation weighting factor and written  $w_R$ . The equivalent dose, written  $H_T$ , in a tissue or organ is given by the following expression:

$$H_T = \sum_R D_{T,R} * w_R \quad (2.3)$$

where  $H_T$  is the equivalent dose for organ T,  $w_R$  is the weighting factor for radiation R,  $D_{T,R}$  is the average absorbed dose in tissue or organ T due to radiation R. ICRP Publication 60 assigns the weighting factors  $w_R$  with each type of radiation, as shown in Table 2.1. Unit for equivalent doses is the Sievert (Sv). The values of  $w_R$  are the result of estimates

**Table 2.1:** Weighting factor  $w_R$  for different types of radiation, ICRP Publication 60.

Radiation	Energy	Values of $w_R$
Photons	-	1
Electrons	-	1
	< 10 keV	5
Neutrons	10 - 100 keV	10
	100 keV - 2 MeV	20
	2 - 20 MeV	10
	> 20 MeV	5
Alpha, fission fragments, heavy nuclei	-	20

conducted by comparing the relative biological effectiveness (RBE) of different radiations in inducing cancer in an organ. According to experiment, values are significant only at low doses of radiation, which lead to stochastic effects[7].

**Effective dose** Epidemiological studies have shown that occurrence of cancers depends on the intrinsic sensitivity of each organ. So, each tissue or organ is associated with a weighting factor  $w_T$  which takes into account the probability of radiation-induced stochastic effects in the organ or tissue. This factor allows calculation of the effective dose  $E$ , a hypothetical dose which, administered uniformly to the entire body, would cause the same late-onset damage as all the doses received by the same individual on the different organs separately at different times, so it makes possible to estimate the risk for humans from a measurable quantity, the absorbed dose. In case of partial exposure of several organs, the effective dose is:

$$E = \sum_T w_T H_T \quad (2.4)$$

in Sievert unit. Values of some tissue are shown in Table 2.2.

**Kerma** In order to give a conceptionally clear description of what could happen when a neutral radiation (gamma or neutron) interacts with matter a special quantity is defined which comprises the total kinetic energy transferred to secondary particles per mass element. It is called *KERMA* (kinetic energy released per mass) and is measured in  $Jkg^{-1}$ . Each mass element is, of course, not isolated but part of its environment to which it loses particles and from where others enter. Secondary particle equilibrium is obtained if every particle leaving the element is compensated by an entering one of exactly the same type and energy. A necessary - but not sufficient - condition for this to occur is that the mass element is part of a homogeneous medium at a depth that is larger than the range of the most energetic particle. It is immediately clear that this can never happen at or near surfaces. Kerma and dose, however, are even in the case of secondary particle equilibrium not generally identical; this is only if *bremstrahlung* losses are negligible. At the surface there is the largest difference because of the lack of equilibrium, see Figure 2.1 . They

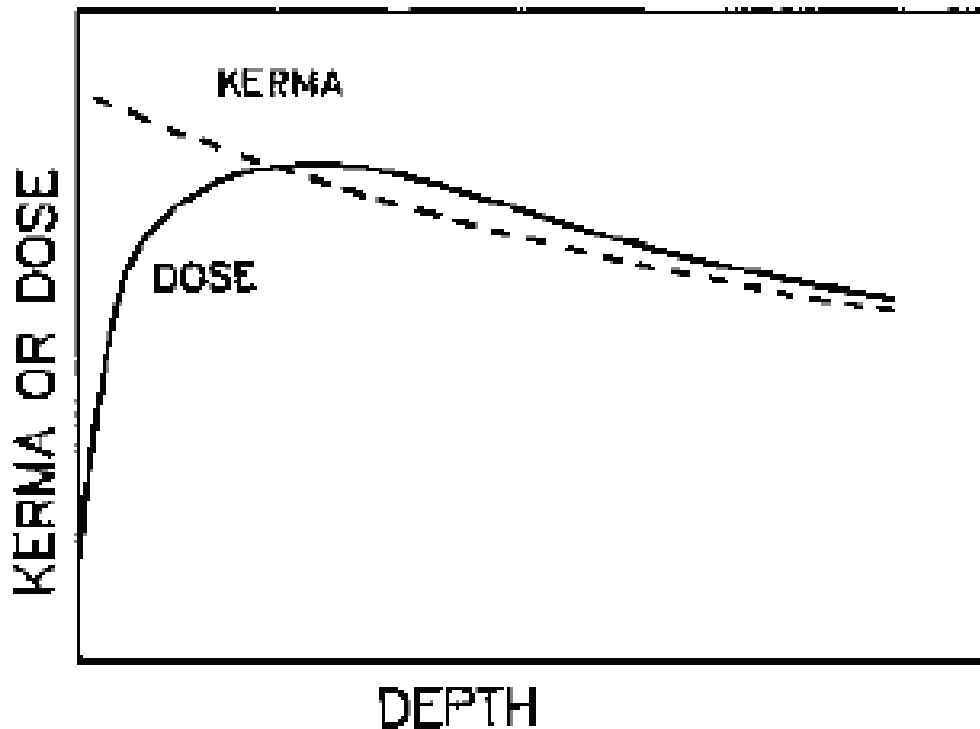


Figure 2.1: *Difference between Kerma and Dose*[9].

become smaller with greater depths where they then are only due to bremsstrahlung.

ii. LET

The energy deposition in an exposed body is mediated almost exclusively by charged particles. These ionize atoms on their way losing parts of their energy in successive steps

until they reach the end of their range. Depending on the type of particle, ionizations are more or less closely spaced, which is very important if one considers energy deposition onto very small sites. This situation may be described by the energy loss of a particle per distance travelled. Corresponding quantity is called *linear energy transfer* (LET) which is defined as the amount of *locally* absorbed energy per unit length. Locally means that energy absorbed is a fraction of total energy transferred in which there is a limit on energy deposited in a local site of medium; 100 eV has been widely accepted, which corresponds to an electron range of about 5 nm. LET without restriction on energy is equal to the stopping power:

$$LET_{\infty} = -\frac{dE}{dx} \quad (2.5)$$

This physical quantity is the first step to evaluate the biological damage. Since the LET value is proportional to the energy transferred by the radiation, it is related to the ionization density and to biological effects[10].

### iii. RBE

For a simple quantitative description of the dependence of radiation quality, the concept of "relative biological effectiveness" (RBE) has been introduced. This parameter is defined as the ratio of doses which yield the same effect if one compares the test radiation with 250 kV X rays or *cobalto* – 60 gamma rays:

$$RBE = \frac{D(250kVXrays)}{D(test \ radiation)} \quad (2.6)$$

RBE shows a dependence on LET. In fact, to a higher LET corresponds a greater number of ionizations along the path inducing a more important damage and this, as mentioned before, is associated with a high RBE value. This parameter is plotted versus  $LET_{\infty}$  in Figure 2.2. The overall behaviour is: RBE increases with LET, passes a maximum around 200 keV/ $\mu$ m from where it declines. This last part has been related to the *overkill* situation where more energy is deposited per particle traversal than actually required for the inactivation of cells. The initial rise is commonly interpreted to mean that essential lesions, like double strand breaks of DNA, are formed with greater efficiency with higher ionization density[12].

**Proton RBE** Proton effects on tissue is still an open question of physics and biology. Since initial studies, dose specification in proton radiation therapy was fixed to a constant RBE value, while after a lot of studies is obtained that other variables influence RBE, like position in the Bragg Peak, initial beam energy and tissue. The physics process that could influence the most is fragmentation. When the incident beam interacts with the target in an inelastic collision, a non-negligible amount of fragments is produced outside of the planned area. They can derive from the target or from the projectile disintegration, according to the different conditions in which the fragmentation takes places (see Section

IV). In case of a proton beam, target nuclei could fragment and change RBE, as is shown in Figure 2.3.

#### iv. DNA damage

In DNA, radiation alterations may be classified in *single strand breaks* and *double strand breaks*:

**Single strand breaks** this is a scission in one chain, while the other remains unaffected.

**Double strand breaks** this is usually caused by a single energy deposition event or by the interaction of two SSB formed individually in close proximity.

DBSs are more likely to happen when heavy ions interact with matter, because ionized electrons, created in this process, have a mean free path of the order of few nanometers. This process cause permanent damage to the DNA because both chains are broken on the same spot and it is not possible to retrieve the genetic information. This means that heavy ions have a higher damage capability than photons, which, instead, are involved in the SSB (Figure 2.4). An indirect effect on DNA is the ionization of molecules such as  $H_2O$ , giving  $H^+$  or  $OH^-$ , that can attack other molecules. For example  $OH^-$  attacks the DNA molecules, which have very soft electrical bounds. Damage depends on the kind of alteration of the DNA, causing cancer or long term genetic mutations.

## II. APPLICATION IN CANCER THERAPY

Treatment with radiation plays an important role. The most used radiations are photons, because of the advanced knowledge on how to produce them and modulate in energy.

### i. Radiotherapy

Radiotherapy uses photon beams with energy between 6 MeV and 25 MeV like in Figure 2.5 and also electron beams in combination with the other therapy. Initially X rays were mainly used but are now substituted with  $\gamma$  rays that are far more energetic. The photon beam used in radiotherapy shows a characteristic depth-dose profile that follows the absorption law and, therefore, displays a steep exponential decrease of dose with depth, like in Figure fig:radio. It presents a maximum between 1 cm and 5 cm. The problem of radiotherapy is that localization of the maximum dose release that, even at greater energies, remains near the surface of the target while tumors are often found in depth[15].

### ii. Hadrontherapy

Hadrontherapy is an oncological technique that uses hadrons beams, not photon beams, accelerated by cyclotrons or synchrotrons to energies from 50 MeV/u to 400 MeV/u (per nucleon). The strenght of hadrontherapy lies in the unique physical and radiobiological



properties of these particles; they can penetrate the tissues with little diffusion and deposit the maximum energy just before stopping. This allows a precise definition of the specific region to be irradiated.

The idea of using protons for cancer treatment was first proposed in 1946 by the physicist Robert Wilson, who later became the founder and first director of the Fermi National Accelerator Laboratory. The first patients were treated in the 1950s in nuclear physics facilities by means of non-dedicated accelerators. In the late 1970s improvements in accelerator technology, coupled with advances in medical imaging and computing, made proton therapy a viable option for routine medical applications.

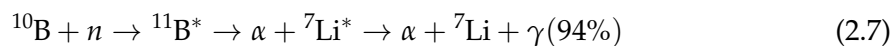
In Italy there are three facilities: CATANA Proton Therapy beam line in Catania, with a energy up to 60 MeV, CNAO in Pavia with a energy up to 250 MeV for protons and 400 MeV/u for carbon beam and APSS in Trento with energy between 60 and 230 MeV for protons.

This therapy shows a distinctive dose release profile that makes it more effective than radiotherapy; it is characterized by a distinct narrow peak, *Bragg peak*, at the end of the particles path with a sharp fall-off at the distal edge (Figure 2.6). Distribution shows a small entrance dose, a well-defined range and a small lateral beam spread.

Despite being both employed in the therapy, protons and carbon ions have different features for what concerns the biological effects and the dose tail. At same absorbed dose, protons have a similar biological response to photons, while heavy ions show higher effectiveness. In addition heavy ions exhibit a distinctive dose tail beyond the Bragg peak, caused by secondary fragments produced in nuclear reactions along the stopping path of ions[16].

### iii. Boron Neutron Capture Therapy

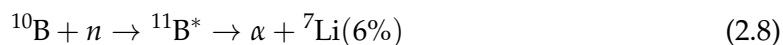
The *Boron Neutron Capture therapy* (BNCT) is a sperimental radiotherapy based on irradiation with termal neutrons on a cancer region enriched with  $^{10}\text{B}$ . Neutron capture reaction with boron lead to formation of  $^{11}\text{B}^*$  excited which decays suddently into two products with an high LET very energetic, an alpha particle and a  $^7\text{Li}$  by means of two nuclear reactions:



where products energies are:

- $E_{\alpha} \simeq 1.47\text{MeV}$
- $E_{^7\text{Li}} \simeq 0.84\text{MeV}$
- $E_{\gamma} \simeq 0.48\text{MeV}$

and



where products energies are:

- $E_{\alpha} \simeq 1.78\text{MeV}$

- $E_{7\text{Li}} \simeq 1.01\text{MeV}$

The products ionize matter near capture region, into  $5\mu\text{m}$  for  $\alpha$  and  $9\mu\text{m}$  for  ${}^7\text{Li}$ , a not much small volume of a cell ( $\sim 10\mu\text{m}$ ). The advantages to use boron are that it is not a radioactive isotope, it is available in nature and very easy to link into molecules, and than products of reaction have high values of LET and small ranges, so they release energy inside cancer cells. Although cross section of capture for other nuclei are much smaller than boron ( $\sigma_{10\text{B}} \simeq 3.84 \cdot 10^3\text{barn}$ ), two of these, hydrogen and nitrogen ( $\sigma_{1\text{H}} \simeq 0.33\text{barn}$ ,  $\sigma_{14\text{N}} \simeq 1.75\text{barn}$ ), are more concentrated than boron and absorb neutron. So the treatment is optimized in order to have a lethal effect on cancer by using  $\sim 10^9$  atoms of  ${}^{10}\text{B}$  for a cell with a thermal neutron fluence of  $10^{10} - 10^{12} \frac{\text{neutrons}}{\text{cm}^2}$  and a boron concentration inside tumor region of  $35 - 50 \frac{\mu\text{g}}{\text{g}}$ . Disadvantages of this therapy are linked to boron distribution, because an ideal drug should fill tumor region and stay outside blood-brain barrier with a small concentration into blood. Another problem is how to control neutron flux from a nuclear reactor and focus it on the tumor region[17]. In figure 2.7 is shown main reaction for BNCT.

#### iv. Proton Boron Capture Therapy

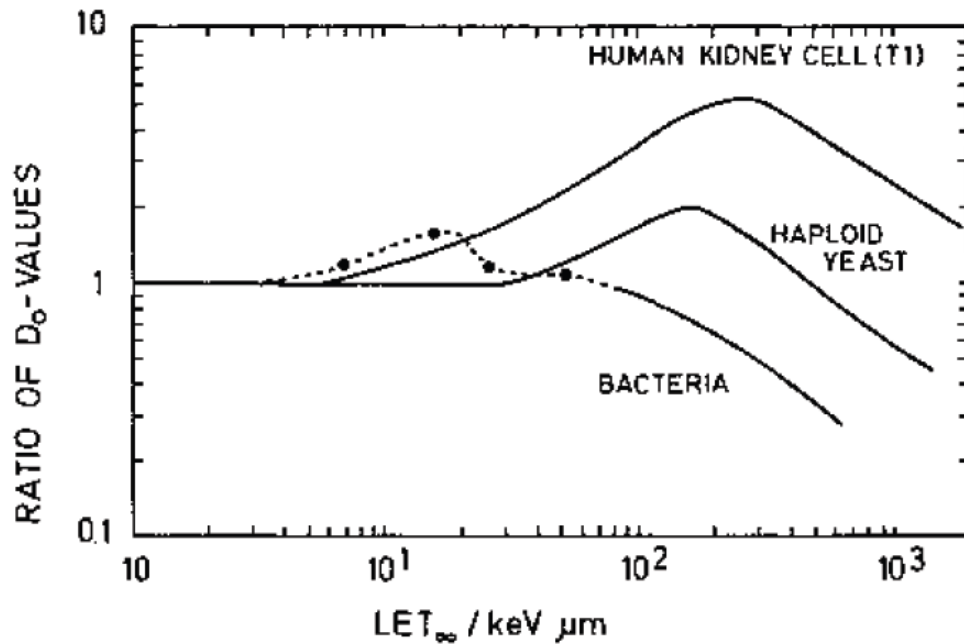
Besides the advantages of using a neutron-free nuclear fusion reaction, the relevance of this method stems from the fact that the  $p + {}^{11}\text{B} \rightarrow 3\alpha$  cross section becomes significantly high at relatively low incident proton energy, for example around the Bragg peak region. As shown in Figure 2.8 a proton beam as conventionally used in protontherapy is drastically slowed down across the tumor (the Bragg peak region). Thus, most of its energy (dose) is delivered to the tumor cells. Under the assumption that a given concentration of  ${}^{11}\text{B}$  nuclei is present preferentially, but not exclusively, in the tumor, the arrival of slow protons could trigger a series of fusion events generating several alpha particles that are localized in the tumor region. In fact, most of the alpha particles generated in the proton-boron reaction have an average range in water of less than  $30\mu\text{m}$ , thus comparable with the typical cell size. Hence, even if such particles are mainly produced outside the cell cytoplasm due to sub-optimal boron uptake, the probability that they would reach the nucleus and damage the DNA remains very high. Moreover, even if a non-negligible concentration of  ${}^{11}\text{B}$  nuclei is present in the healthy tissues surrounding the tumor, the number of generated alpha particles, would be relatively low, or completely absent, due the non-favourable incident proton energy spectrum away from the tumor region. This would lead to a more biologically effective particle dose localization, higher than the one currently achievable with conventional protontherapy, thus to a more efficient treatment in terms of an *enhancement* in cancer cell lethality, especially because of the clustered nature of the DNA damage, caused by the high-LET alpha particles emitted in the tumor region. Hence, protontherapy could acquire the benefits of an enhanced efficiency in cancer cell killing, moving close to  ${}^{12}\text{C}$  ion hadrontherapy but without the above-mentioned complications of the latter.

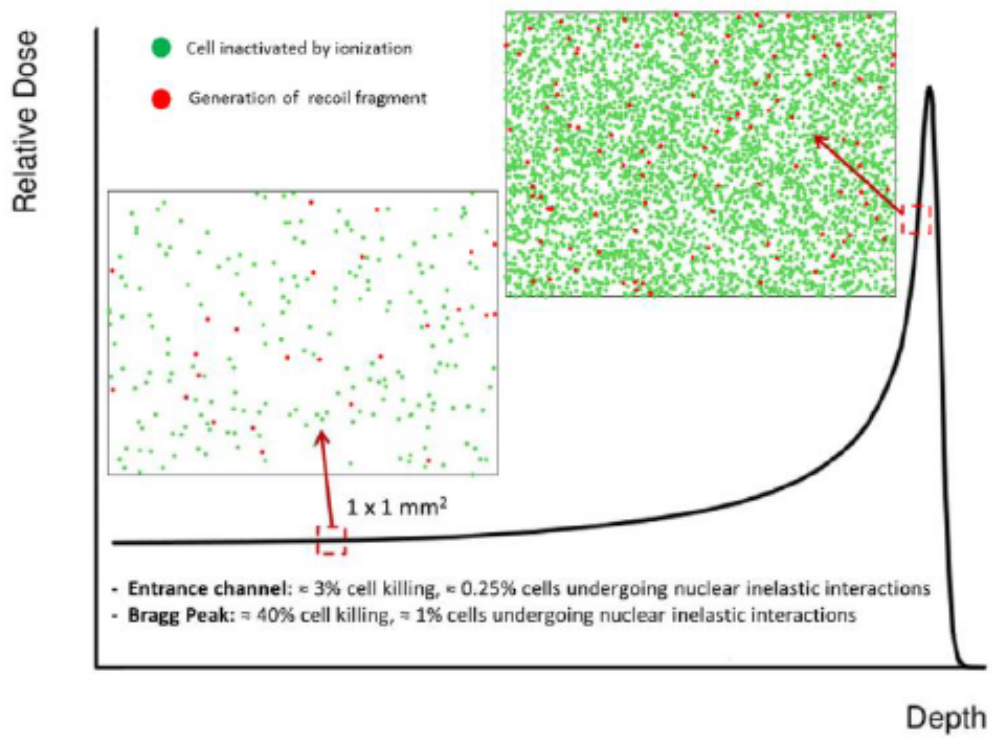
Ballistic advantage granted by the inverted dose-depth profile of charged particles is

such that in protontherapy most of the dose is released mainly in the tumor region (left panel). If cancer cells are loaded with  $^{11}\text{B}$ -delivering agents, unreparable DNA clustered lesions will be also produced by High-LET alpha particles. For a given clinical case, such higher *dose modifying factor* can potentially allow to reduce the overall dose delivered to the patient compared to a standard treatment without the presence of  $^{11}\text{B}$ -delivering agents in the tumor.

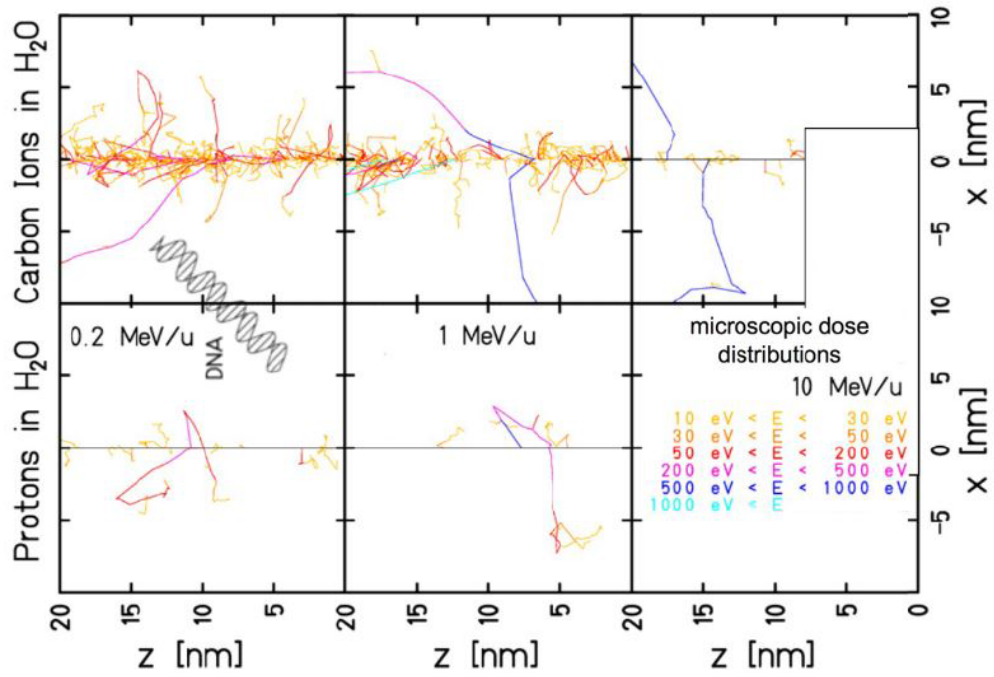
**Table 2.2:** Weighting factor  $w_T$  for different types of tissue [8].

Tissue	Values of $w_T$
Gonads	0.08
Bone marrow	0.12
Colon	0.12
Lung	0.12
Stomach	0.12
Bladder	0.04
Breast	0.12
Live	0.04
Oesophagus	0.12
Thyroid	0.04
Skin	0.01
Bone surface	0.01
Brain	0.01
Salivar glands	0.01
Other tissues or organs	0.12

**Figure 2.2:** Relative radiation sensitivity of various cell types as a function of LET[11].



**Figure 2.3:** Cell inactivated by ionization (green) and target fragmentation (red) in tissue of 1 mm<sup>2</sup> [13].



**Figure 2.4:** Simulation with the Montecarlo code of the interaction between carbon ions and protons with matter in proximity of the range end, where particles slow down and their energy is of few MeV per nucleon. The graph shows the tracks of single secondary electrons. Carbon ions produce more tracks than protons, increasing the probability of having direct damage to the DNA[14].

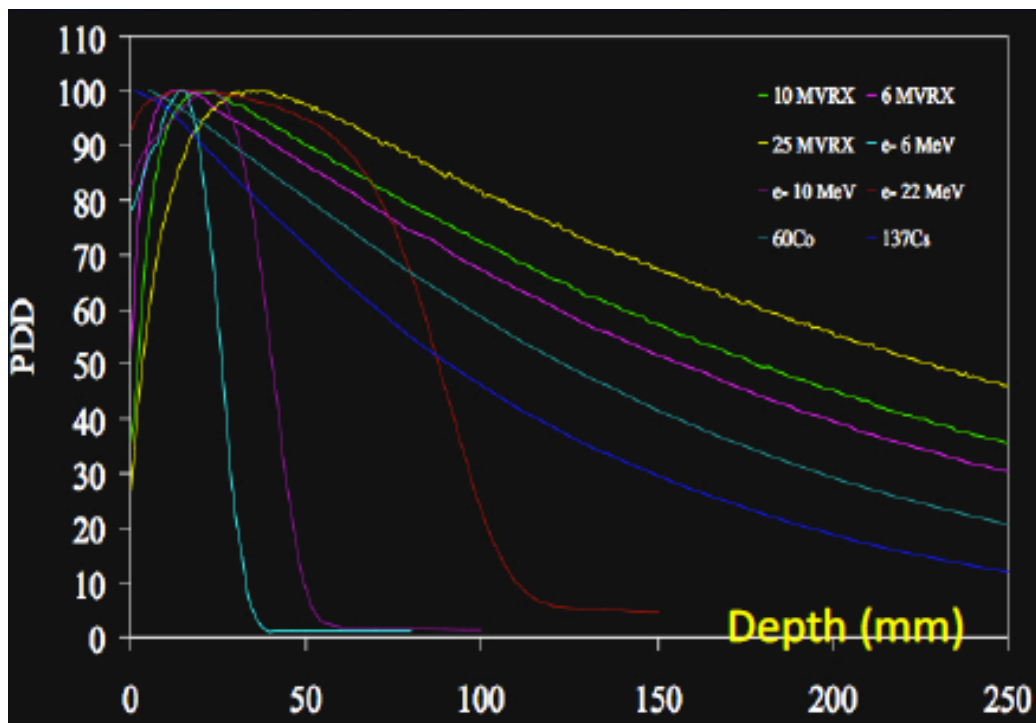


Figure 2.5: Percent Depth Dose of photons beams for several photon energy.

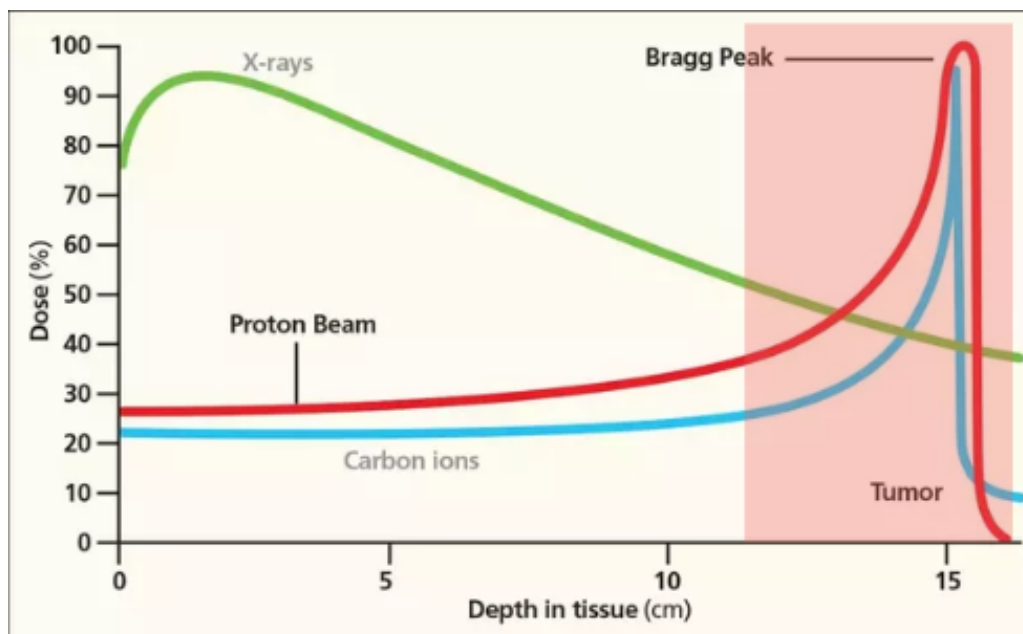


Figure 2.6: Comparison of depth-dose profiles in water of photons, protons and high-energy carbon ions.

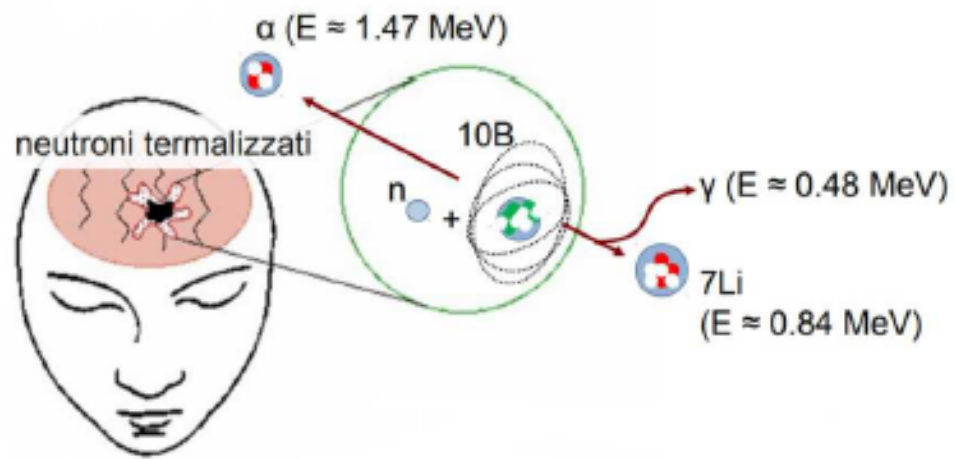
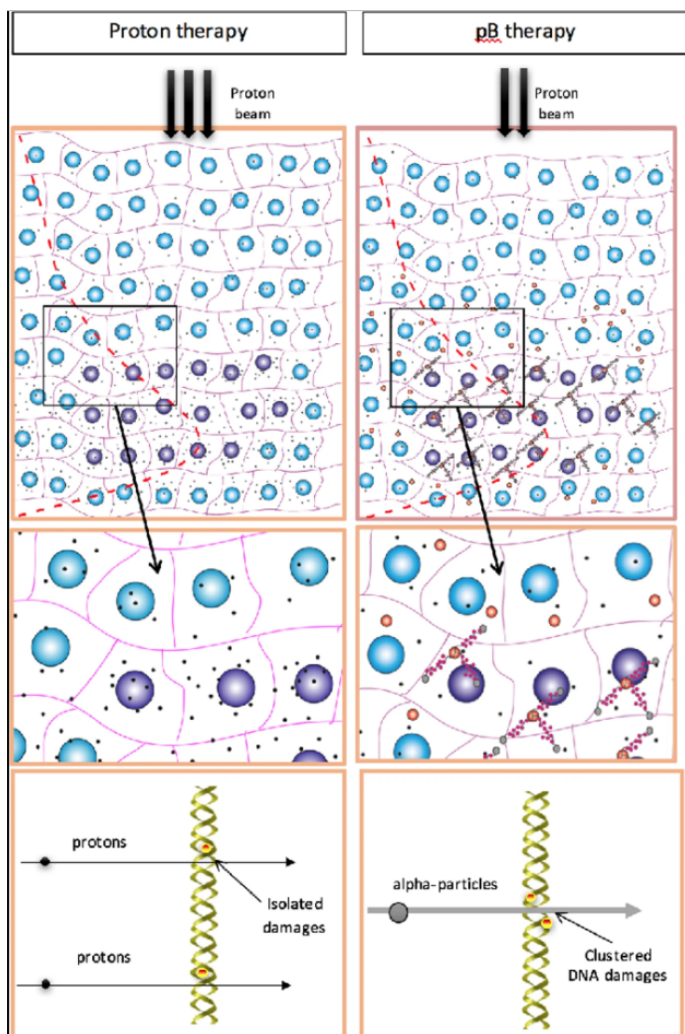


Figure 2.7: Rappresentation of BNCT.





**Figure 2.8:** Representation of PBCT: left panel shows conventional protontherapy, in which incident proton beam mainly results in isolated, repairable DNA breaks; right panel shows how proton-boron extremely localized alpha emission in the Bragg peak causes irreparable clustered DNA damage [18].

## Chapter 3

# ERHA project

The technology of proton's production and acceleration is a part of several research projects. The ERHA is focused on the realisation of an hadrontherapy center, in which the accelerator is linear and compact. The main reason is the possibility to install this machine inside a hospital centre. In order to build this type of accelerator, there are some specification to be setted: type of particle, for example protons, but also ions, energy of the beam, number of particle accelerated in a second, and repetition frequency. In all the situation accelerator is placed inside a  $100\text{ m}^2$  room, with all the instrumentation.

### I. LINAC ACCELERATOR

LINAC is a type of particle accelerator that is based on the multiple acceleration method: in the gap between two beam pipe tubes there is an electric potential difference  $V = V_0 \cos \omega t$  that establishes a variable electric field, so the particle is accelerated if it crosses gaps in a time synchronous with field, so for every wave period. After a period, particle gains velocity, so the distance between gaps has to increase, roughly as  $L = vT/2$ , where  $v$  is the particle velocity[19]. ERHA project is based on some working conditions which are shown in Table 3.1 Some characteristics are described in a simplified way:

- **Peak current:** Mean current, that correspond to mean number of particles for a treatment, is  $1 \div 10\text{ nA}$ . A pulsed beam with a repetition rate of  $10\text{ Hz}$  and with a pulse width of  $4\mu\text{s}$ , has a peak current of:

$$\frac{1\text{nA}}{10\text{Hz} \cdot 4\mu\text{s}} = 25\mu\text{A} \quad (3.1)$$

- **Number of protons per injector pulse:** Peak current of the injector is  $1.0\text{ mA}$ . If pulse width is  $4\mu\text{s}$ , number of protons is:

$$\frac{1\text{mA} * 4\mu\text{s}}{1,6 * 10^{-19}\text{C}} = 2,5 * 10^{10}\text{ protons} \quad (3.2)$$

per pulse, while per second are  $2,5 * 10^{10} * 10\text{Hz} = 2,5 * 10^{11}\text{ p/s}$

- **Approximative lenght:** A general solution for lenght of an accelerator is:

$$L = \frac{T_{final}}{qE_{0T} \cos \phi_s} \quad (3.3)$$

where  $q$  is the charge of proton,  $E_{0T}$  is the effective accelerating field of  $10 \div 15$  MV/m and  $\phi_s$  is the sincronous phase of  $-20^\circ$ . Typical lenght is 20 meters.

- **RF Power:** In order to accelerat it needs a radio frequency power at peak:

$$P_p = \frac{(E_{0T})^2}{Z_{eff}} L \quad (3.4)$$

where  $Z_{eff}$  is the shunt resistance for unit lenght. So  $P_p$  is some MW. In order to obtain mean power,  $P_p$  must be multiplied by duty cycle.

- **Accelerator modules:** Protons are initially accelerated by the Radiofrequency Quadrupole (RFQ) to 4 MeV. For energies from 5 MeV and 100 MeV are usually used Drift Tube Linac modules (DTL), working at less than 500 MHz. After these energies are used Side Couple Linac modules (SCL), working at 3 GHz.
- **Frequency:** In general accelerators work at high frequency, because it reduces dimensions in which is located, but also RF power, and break-down limit is higher, as obtained by Kilpatrick law  $E_{break} \propto f^{0.4}$ .

### i. Injector

The first module for injecting protons into accelerating pipe is the Accsys-Hitachi PL-7, a commercial model made of a duoplasmatron source of 30 keV and a RFQ of 4 MeV operating at a 428,27 MHz, seventh subarmonic of 2997,92 MHz, the operating frequency of the LINAC. The power supply consists of 15 planar triods, EIMAC tubes (CPI-YU176A), 1 for pre-amplificator, 2 for second pre-amplificator and 12 for the final amplificator[20].

**Duoplasmatron** Proton source is a ions source supplied with gas, typically hydrogen. A schematic figure is 3.1. An electric arc, with low pressure, between anode and cathode is produced. On cathode there is a thin wire, typically made of tungsten, in which flows current which produces electrons by thermoionic effect. On cathode flows gas which is ionized by electric arc and produced plasma, which is confined by an intermediate electrode conic shaped and stay in little region *A*. Although plasma is globally neutral, particles inside are charged and let it expand. In order to confine plasma, it is used a magnetic field that mantain a pressure on plasma. On electrode there is a little hole that made particle flowing to anode. Between intermediate electrode and anode there is an other electrode that is positive and pick electrons. In region *B* protons are accelerated towards extraction electrode and goes into a region with a lense and a diafram, which are used to focus the beam[21].

**RFQ** The RFQ is a linear accelerator that accelerates, focuses and make bunches of protons with a radiofrequency field. It is the best way to accelerate ions at low energies, and it is simple to design and build, and accelerate intense beams with 90% of efficiency. RFQ structure consists of a radiofrequency cavity made by four electrodes (Figure 3.2) formed and shaped properly in order to obtain right acceleration and focusing. Protons are focused polarizing electrodes symmetrically, in order to create a quadrupole field. Protons are simultaneously accelerated by the same field by shaping electrodes longitudinally like a sinusoid function, in Figure 3.3. Peaks of electrodes are separated by a length of  $\beta\lambda$ , in order to shape bunches of protons[22].

## ii. SCDTL

The end of injector is matched with an ERHA project developed module which is based on a Drift Tube Linac in which accelerating tanks are alternated to pairing tanks situated off axis, in a way that is possible to pair fields inside tanks. Between two accelerating tanks there are quadrupole permanent magnets in a FODO cell, that focus the beam. This is an innovative module, because it is designed to be modular and easy to set, see Figure 3.4 that shows the Module 0, an SCDTL (*Side Coupled Drift Tube Linac*). A schematic visualization of how SCDTL works is shown in Figure 3.5 where are visible tanks that consist of a series of drift tube cell, from 3 to 7  $\beta\lambda$  long, while the space between tanks is used to focus the beam by means of a quadrupole (PMQ) which is 3 cm long and with a diameter 2 cm long. The Drift Tube Linac is made by an array of resonant cavities, at RF frequency. The synchronization of the particle motion with electromagnetic oscillations is made by changing length of drift tubes between gaps. The length increasing of the cavities increases also the capacity, and decreases resonance frequency. In order to keep frequency to a constant value, gap geometry in tanks have to be shaped reducing drift tube diameters progressively, like in Figure 3.6. Design of cavities creates also the right distribution of the field and establishes the parasitic currents on the separation surface between two cavities. An electromagnetic wave that is resonant in a cavity is depicted by a *mode*, that could be *Transverse Electric* (TE) if the components of the field that is directed along propagation direction is  $\vec{B}$ , otherwise *Transverse Magnetic* (TM), in order to satisfy contour conditions where field is null. Electric field distribution and parasitic currents for the  $TM_{010}$  are in a shape that leaves field distribution equal when the separation surfaces are eliminated. In order to hold drift tubes there are sustain bars in a position in which field is radial and is not perturbed.

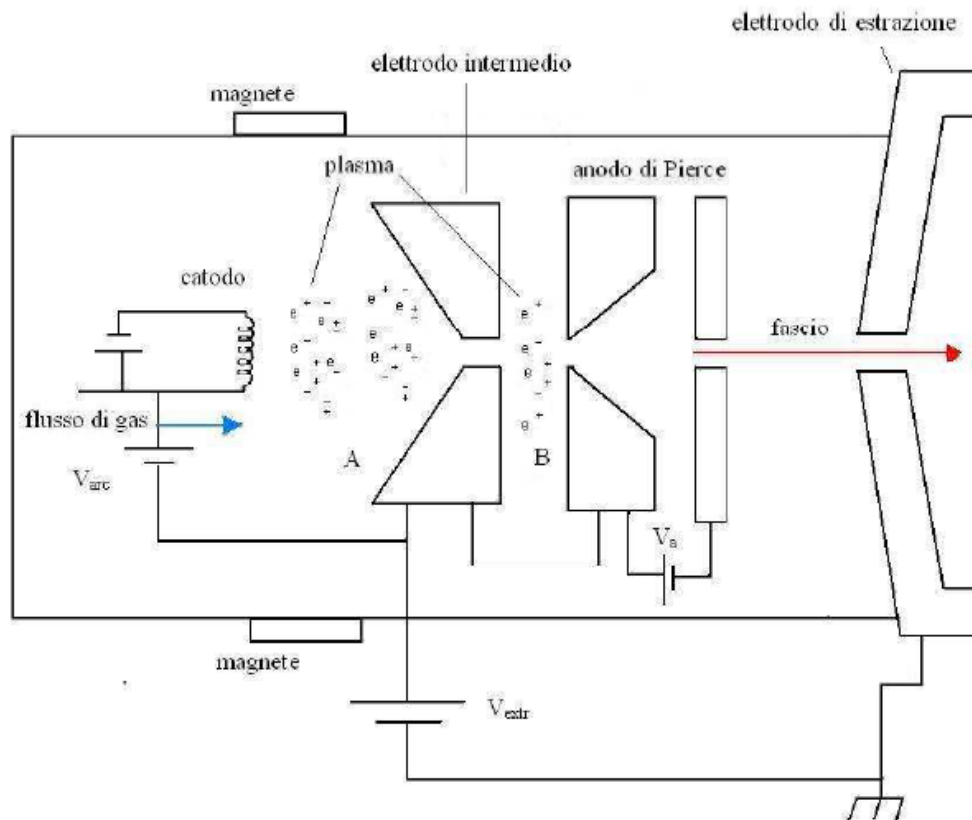
A big problem on DTL is to sustain the mode  $TM_{010}$ . In a complex structure is a very common problem that different resonant modes are generated which could give rise to significantly beam losses. In order to resolve it, some tuning elements are inserted in the structure, so called *post couplers*, orthogonal to sustain bars[23].

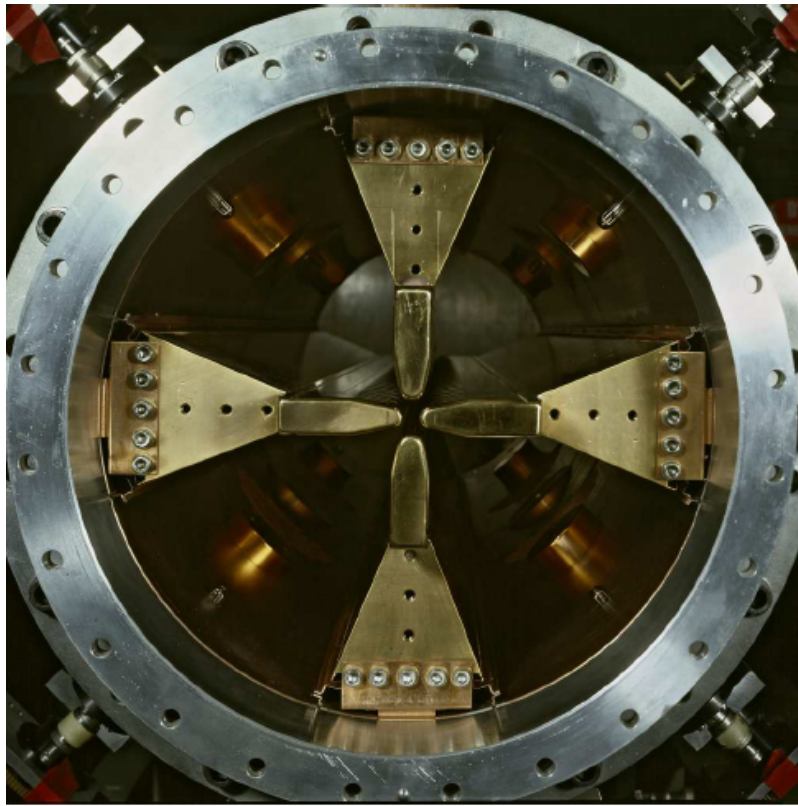
## iii. CCL

CCL module is developed at Los Alamos Laboratory. The structure is similar to SCDTL, it is compact, every cell is  $\beta\lambda/2$  long assembled in tanks of limited length (in a tank there are 16 cells). Between tanks are placed quadrupoles, every tank is made by  $n$  accelerating cell and  $n-1$  pairing cell in  $\pi/2$  field mode, while a bridge coupler pairs the two tanks (see Figure 3.8). An efficient gain of the structure for small energies (bigger than 25-30 MeV) is possible with a "PALME" structure, much more complex to design, as shown in Figure 3.7 High values of the field  $E_0$  can be set to obtain smaller accelerator, but RF power increases as  $E_0^2$ . Kilpatrick's limit of break-down  $E_k$  is of  $47MV/m$  for a 3GHz frequency, with a superficial field  $E_s \leq 2E_k$ . For a geometry reason,  $E_s \simeq 6E_0$ , for a value of  $15MV/m$ . When the field is fixed, an algorithm SUPERFISH calculates geometry of the cells. These are assembled in tank by another program, which calculates energy gain for every tank and focusing force of the quadrupole. At the end a simulation program PARMILA, in which are simulated hundreds of particles, can simulate trajectory inside beam-pipe. A CCL, *coupled cavity linac*, is made by resonant cavities coupled by a hole in the structure. They are used to accelerate protons in a range  $0.4 < \beta < 1.0$ . Single cavity is called cell, in an excited mode very similar to  $TM_{010}$ . Coupling is generated by a hole like in Figure 3.9 where in (a) is shown an electric or capacitive coupling due to a higher electric field than magnetic component, while in (b) is shown a magnetic or inductive coupling, due to a higher magnetic field[25].

**Table 3.1:** Initial LINAC specifications.

Proton Spot Size	2 mm
RF frequency	3 GHz
Mean beam current	$1 \div 10$ nA
Peak current (50 Hz)	$0.7 \div 33$ $\mu$ A
Injector peak current	1.2 mA
Repetition rate	$20 \div 200$ Hz
Pulse duration	2 -20 $\mu$ s
Energy precision	$\pm 0.2$ MeV
Water range	$0.1 \div 3.3$ g/cm <sup>2</sup>
Distal decrease dose (80% - 20%)	< 2 mm
Normalized transverse emittance	< $2 \pi$ mm mrad
Duty cycle	0.01%

**Figure 3.1:** Proton source duoplasmatron.



**Figure 3.2:** *RFQ structure.*

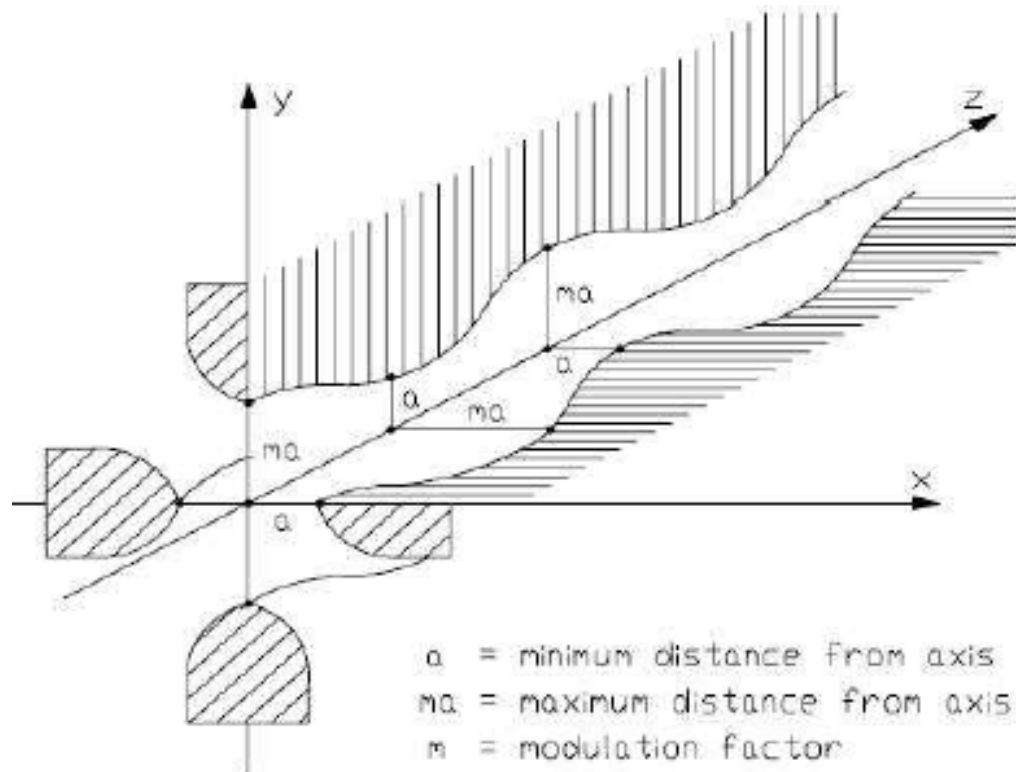


Figure 3.3: Longitudinal modulation of electrodes.

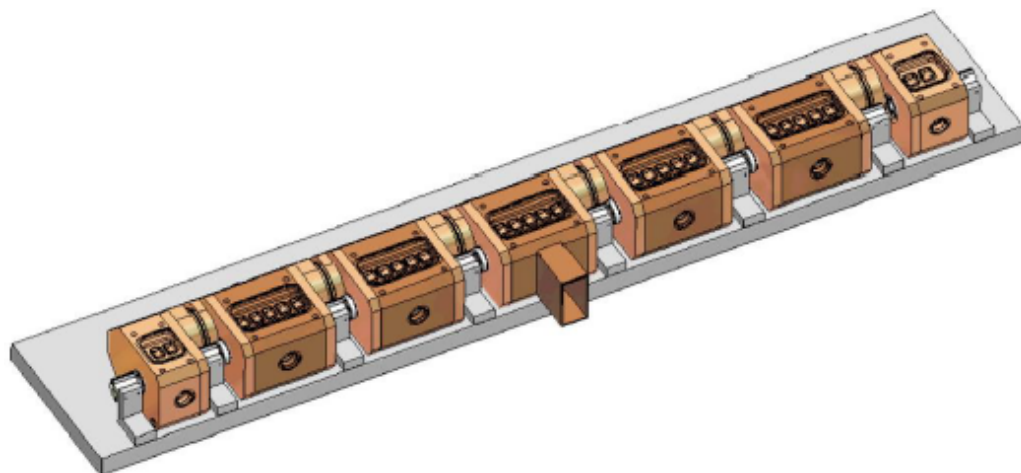


Figure 3.4: Module 0, SCDTL.



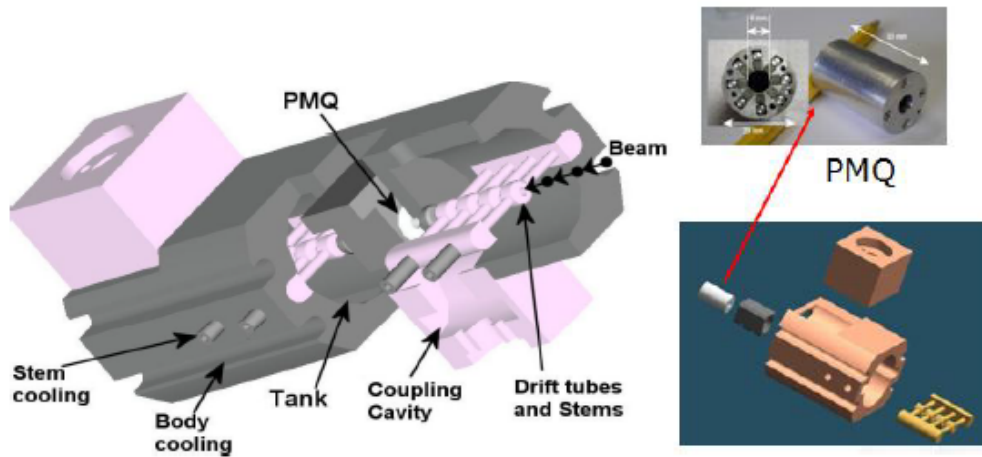


Figure 3.5: Interior part of tanks and PMQ.

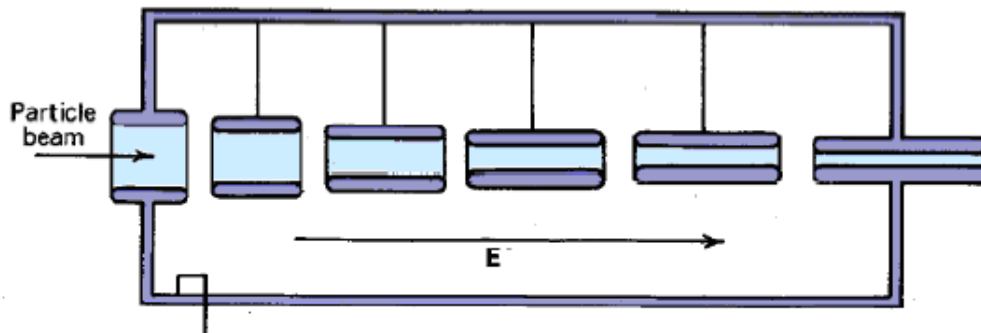


Figure 3.6: Progressive shaped tank.

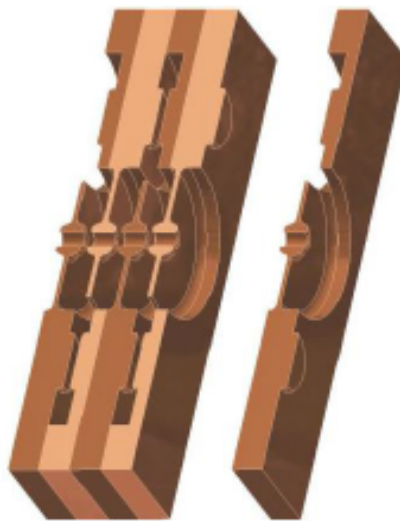


Figure 3.7: PALME structure of CCL.

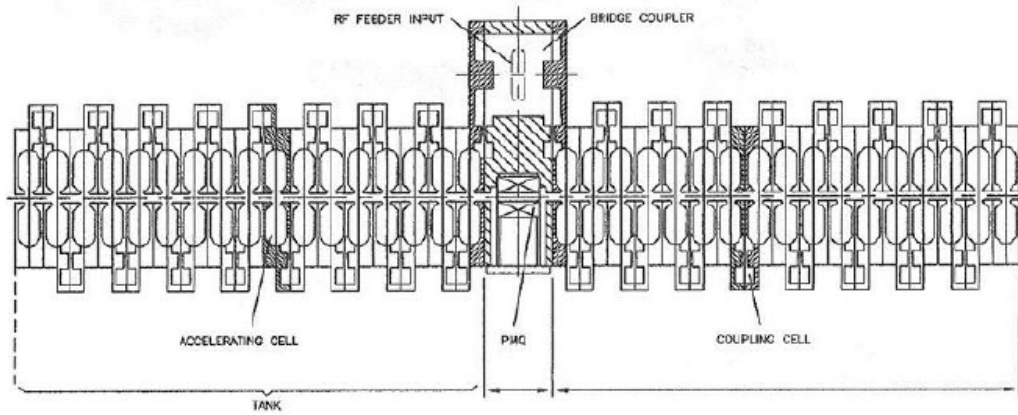


Figure 3.8: [24] Assembled tanks in CCL.

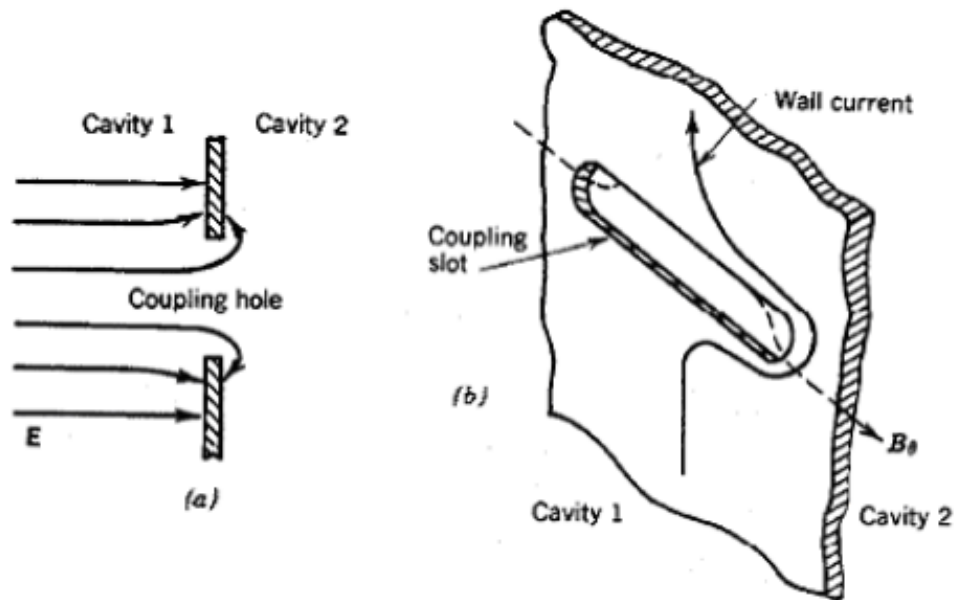


Figure 3.9: Coupling between cavities, electric to left, magnetic to right.



## Chapter 4

# Radiation shielding

The radiological protection aspects of an accelerators are extremely important in the design of these machines. There are many parameters by which particle accelerators may be classified. For example, they may be classified in terms of the acceleration technology, such as power source or acceleration path geometry. Also they may be classified by their application, by types of particle, maximal energy, maximal intensity and duty factor of the beams. In this chapter is described the main radiation source in the bunker and the protection aspects.

### I. PROMPT RADIATION

A basic knowledge of the nuclear reaction mechanisms that happens in the energy range of 1 - 10 MeV is required in order to understand shielding of an accelerator. When a proton hits matter it could undergo a nuclear reaction and form a compound state. The process could give rise to an inelastic scattering, in which the compound could change number of protons and neutrons but  $A$  doesn't change, a so called *charge-exchange reaction*. When  $A$  changes, there is a transfer reaction (*stripping* or *pick-up*) or a *knockout reaction*. The angular distribution of the particle is characteristically anisotropic, peaking in the forward direction. In this situation, each nucleon will undergo further collisions, gradually spreading its excitation energy over the whole nucleus. For a certain time (during pre-equilibrium phase) nuclear state will become increasingly complex, but after a certain relaxation time, statistical equilibrium will be reached. A fraction of the mixture of nuclear states consists of configurations in which energy is concentrated on one nucleon so that it may escape from the nucleus. Similarly, kinetic energy may be concentrated on groups of particles and lead to the emission of  $\alpha$  particles, tritons, deuteron etc. This is similar to evaporation and may be characterized by a nuclear temperature  $\Theta \approx 2 - 8$  MeV, so that spectrum of the emitted neutrons may be described by the following Maxwellian distribution:

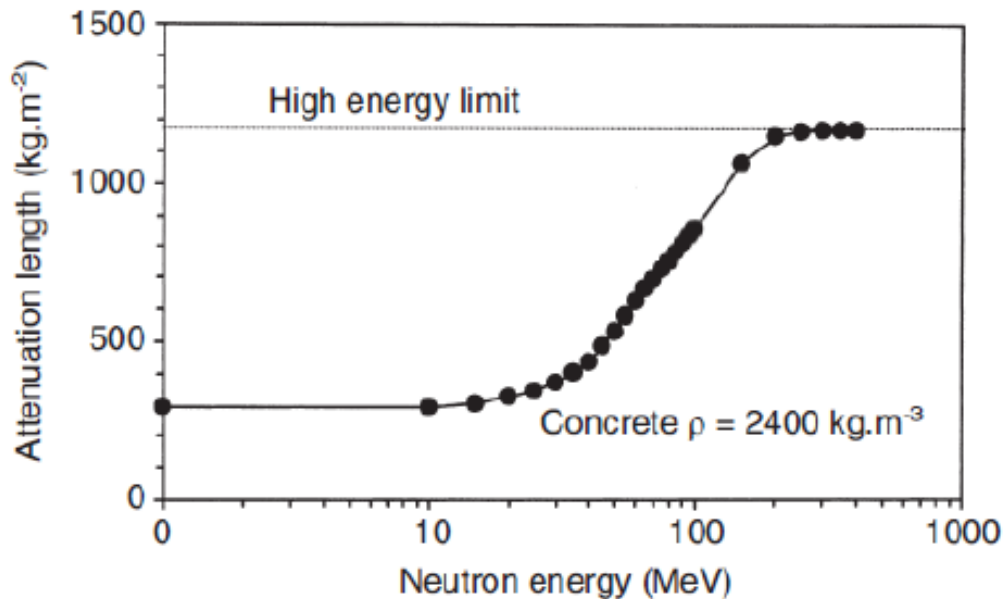
$$dN(E_n) \propto \frac{E_n}{\Theta^2} \exp\left(\frac{-E_n}{\Theta}\right) dE_n \quad (4.1)$$

Compound reactions may occur during the *pre-equilibrium* phase, before statistical equilibrium is achieved. In such cases the angle of emission may still be correlated with the direction of the incident particle. On the other hand, once statistical equilibrium was obtained, the emitted particles have no memory of incident particle and angular distribution is isotropic. The particles emitted could participate in similar reactions resulting in an intranuclear cascade, which develops through interaction of individual nucleons inside nucleus.

Prompt radiation field near interaction point of accelerated protons with matter is complex and becomes more complex as energy is increased. The field consists of a mixture of *charged and neutral particles and photons*. Several simulation codes are available which include all the interaction described above and which allow estimates of the radiation field, such as GEANT4 and Fluka.

In order to attenuate prompt field, in particular as seen above neutron field, two criteria must be satisfied: interpose sufficient mass between the source and the field point and attenuate effectively neutrons of all energies. First criterion is easily obtained by dense material of high atomic mass, while second is most easily met by hydrogen, which attenuates neutrons of all energies via elastic scattering. The two criteria are met by concrete because it contains hydrogen in water form.

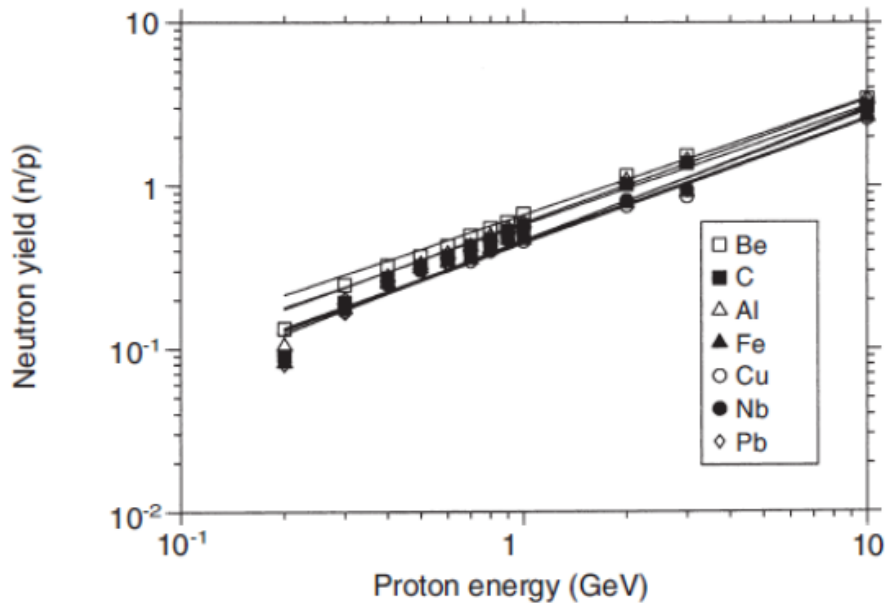
In Figure 4.1 is shown variation of the mass attenuation length,  $\rho\lambda$  for monoenergetic



**Figure 4.1:** Attenuation length  $\rho\lambda$  of monoenergetic neutrons in concrete of  $\rho = 2400 \text{ kg/m}^3$  in function of energy.

neutrons in concrete as function of energy. Below 20 MeV,  $\rho\lambda$  is  $200 \text{ kg/m}^2$ . Above this energy neutron cross section changes from a interaction with target nuclei as a whole, so an elastic scattering, to an interaction with nucleons.

The lower energy neutrons and charged particles are regenerated at all depths in the shield by the inelastic interactions of the neutrons with the shielding material. In other words, at any field point outside the shielding, the highest energy neutrons will be those that have come directly from the source without interaction, or that have undergone only elastic scattering or direct inelastic scattering with little loss of energy and only small angular deflection. Any low energy neutrons and charged particles detected outside the shielding will have been generated by the intranuclear cascade near the outer surface of the shield. In Figure 4.2 the neutron yield is normalized per interacting proton and has a



**Figure 4.2:** The yield of neutrons with energy  $E_n > 100$  MeV per interacting proton in stopping targets of a number of materials as a function of proton energy. The points are the results of calculations with the FLUKA Monte Carlo code and the lines are best fits to these points to the relation  $n(E_p) = n_0 E_p^m$ , [26].

dependence on the proton energy of the form:

$$n(E_p) = n_0 E_p^m \quad (4.2)$$

#### i. Operational quantities

Body-related protection quantities, effective dose, equivalent dose, have drawback of not being amenable to measurement. To meet the requirements of the organizations charged with monitoring workforce exposures, the concept of *operational quantities* was introduced, used to arrive at reasonable assessments of the protection quantities. These operational quantities involve the following characteristic features: they are based on dose at depths of 10 mm and 0,07 mm, respectively, as measured in the ICRU sphere, or in the human body. The ICRU sphere is a reference sphere, 30 cm in diameter, made of tissue-equivalent

material with a density of  $1 \text{ gcm}^{-3}$ . They can be measured at the workplace, by means of external radiation detectors (rate meters, dosimeters), and may be used for individual ambient monitoring purposes at workstations. They serve as estimators, yielding as a rule overestimates ("conservative" estimates), for the effective dose, and the organ-related equivalent doses. When a variety of radiations, energies, and angles of incidence are involved, the respective quantities relating to each of these are additive.

Three major operational quantities are used: two are used for area, or ambient monitoring purposes:

- ambient dose equivalent  $H^*(d)$
- directional dose equivalent  $H'(d, \Omega)$

and third one is used for individual monitoring purposes:

- personal dose equivalent  $H_p(d)$

These quantities correspond to the dose equivalent produced at a point located at a depth  $d$  in a phantom (e.g. the ICRU sphere), or in the human body, which in turn depends on the energy of the radiation involved, and the geometric conditions pertaining to the exposure (direction of irradiation). For routine monitoring purposes, the values found for these operational quantities are deemed to provide adequately accurate estimates for the effective dose, and for the dose to the skin, especially if they are lower than the radiological protection limits.

**Ambient dose equivalent**  $H^*(d)$  is the reference quantity for strongly penetrating radiation, for ambient monitoring purposes.  $H^*(d)$  provides a good estimator for the *effective dose*. As the recommended depth  $d$ , in that case, is 10 mm, this quantity may then be noted  $H^*(10)$ . Many detectors used as dose-rate meters are calibrated with reference to  $H^*(10)$ .

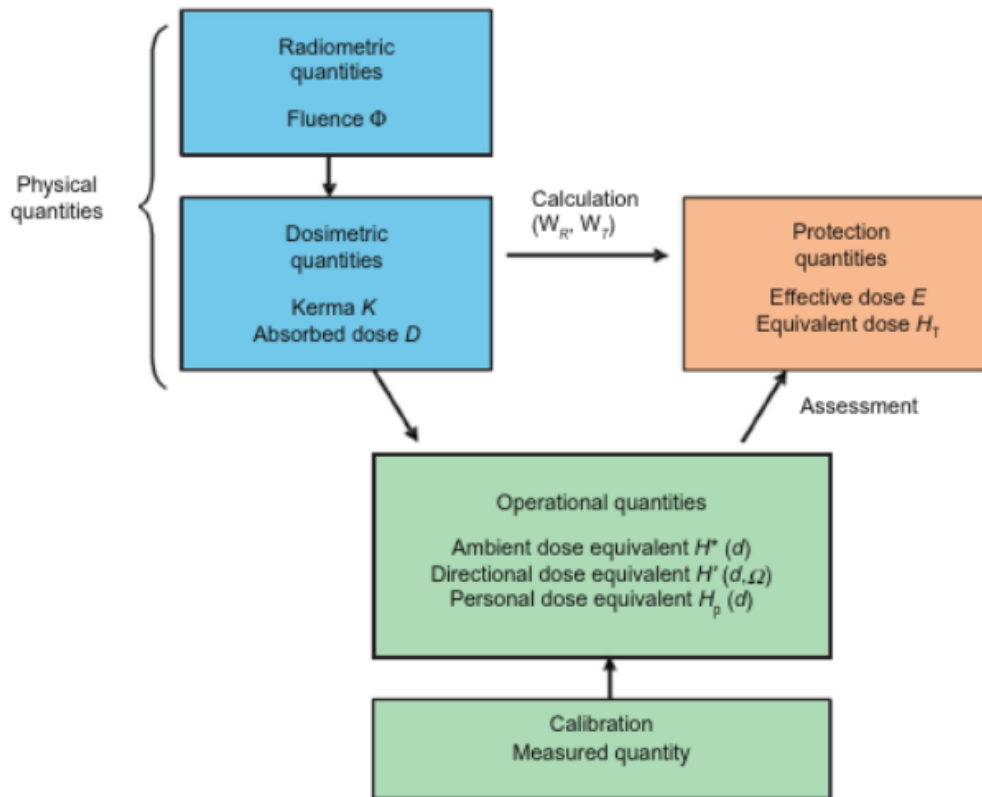
**Directional dose equivalent**  $H'(d, \Omega)$  is the quantity used for low-penetrating radiation, for ambient monitoring purposes.  $H'(d, \Omega)$  stands as an estimator for the equivalent dose to the skin  $H_{skin}$ . Consequently, the recommended depth, in this case, is 0,07 mm. This quantity may thus be noted  $H'(0.07, \Omega)$ .

**Personal dose equivalent**  $H_p(d)$  is the quantity used for personal monitoring purposes. Two cases may arise: for strongly penetrating radiation, the recommended depth stands equal to 0.07 mm, the quantity is then noted  $H_p(10)$ , this providing a good estimator for the effective dose. For low-penetrating radiation, the recommended depth stands equal to 0.07 mm, the quantity is then noted  $H_p(0.07)$ , which is a good estimator for the equivalent dose to the skin  $H_{skin}$ .

Dosimeters worn on the surface of the body, serving for workforce monitoring purposes, are calibrated with reference to  $H_p(10)$  and  $H_p(0.07)$ , they thus yield good estimates for the effective dose, and the equivalent dose to the skin. Such dosimeters are, as a rule,

covered with a tissue-equivalent material.

The depth  $d$ , or indeed any thickness of any given material may be stated in terms of the corresponding density thickness, expressed in  $gcm^{-2}$ , or in  $mgcm^{-2}$ . There are also detectors used for individual monitoring purposes, serving to measure  $H_p(3)$ , this being an estimator for the equivalent dose to the lens of the eye  $H_{lens}$ . In Figure 4.3 is summa-



**Figure 4.3:** Relationships between physical quantities, protection quantities, and operational quantities [7].

rized the relationships between physical quantities, protection quantities and operational quantities.

## ii. Gamma Shielding

Attenuation of photons by various absorbing materials under ideal narrow-beam conditions satisfy the relationship

$$I(x) = I_0 e^{-\mu x} \quad (4.3)$$

where  $I_0$  is the initial photon intensity, a fluence or a flux,  $I(x)$  is the photon intensity after passing through an absorber of thickness  $x$  in narrow-beam geometry, and  $\mu(cm^{-1})$  is the total attenuation coefficient, which accounts for all interaction processes, including scattering reactions, that remove photons from the beam. The attenuation coefficient  $\mu$  is



dependent on the particular absorber medium and the photon energy.

Values of  $\mu$  generally increase as the  $Z$  of the absorber increases because photoelectric interactions are increased in high- $Z$  materials especially for low-energy photons, and high- $Z$  materials yield increases in pair production interactions for high-energy photons. Because of the high- $Z$  effect, lead is often used to line the walls of x-ray rooms, made into lead aprons for personnel protection, and incorporated into leaded glass, and  $\text{BaSO}_4$  is incorporated into concrete (called barite or barytes concrete) to increase its effectiveness as a photon shield.

*Photon attenuation coefficients* in various materials can be calculated from measurements of the intensity of a narrow beam of photons of a given energy. Many calculations of radiation exposure/dose from photon sources are straightforward once the flux is known. A useful formulation is the flux at a distance  $r$  from an attenuated point source:

$$\phi(x) = \phi_0 \frac{e^{-\mu x}}{4\pi r^2} \quad (4.4)$$

The expression  $e^{-\mu x}/4\pi r^2$  is referred to as the *point kernel* which is the response at a point  $r$  from a source of unit strength. The point kernel is used extensively in developing relationships between flux and exposure for various source geometries and absorbing media.

Although many radiation sources can, with its ease and utility, be represented as a point or an approximates a line source, a contaminated area that is representative of a disc or infinite planar source, and various volume sources. Practical approaches can be used to determine the photon flux from point kernels spread over such geometries, and once the flux has been determined it than be applied in the usual way to calculate radiation exposure [27].

### iii. Neutron Shielding

The interactions that slow neutrons down and cause their eventual removal from a beam are probabilistic: they either occur or they do not. Consequently, a flux of neutrons of intensity  $I$  will be diminished in a thickness  $x$  of absorber proportional to the intensity of the neutron source and the neutron removal coefficient  $\Sigma_{nr}$  of the absorbing material:

$$-\frac{dI}{dx} = \Sigma_{nr} I \quad (4.5)$$

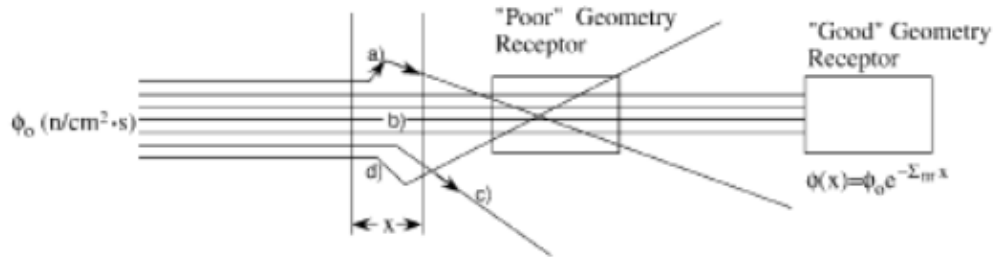
which has a solution

$$I(x) = I_0 e^{-\Sigma_{nr} x} \quad (4.6)$$

like a photon attenuation process, where  $I_0$  is the initial intensity and  $I(x)$  refers to those neutrons that penetrate a distance  $x$  in an absorber without a collision. Therefore  $e^{-\Sigma_{nr} x}$  represents the probability that a given neutron travels a distance  $x$  without an interaction. Conceptually,  $\Sigma_{nr}$  can be thought of as the probability per unit path length that a neutron will undergo an interaction as it moves through an absorber and be removed from the

beam either by absorption or scattering. In this context then it very much resembles the attenuation coefficient for photons in *good (narrow-beam) geometry*, and can be similarly developed and used for neutron shielding and dosimetry.

The features of neutron beams, including the concept of *narrow-beam* effects, are shown in Figure 4.4, where different interactions remove a neutron which doesn't reach the detector.



**Figure 4.4:** Absorber of thickness  $x$  that depletes a beam of neutron because of (a) an inelastic scattering followed by a photon emission and a scattering reaction back into beam, or (b) an absorption or capture (grey lines), (c) elastic scattering out of the beam, or (d) elastic scattering with additional scattering back into the beam.

As a consequence of the complicated resonance structure of the neutron cross-section, neutron removal coefficient  $\Sigma_{nr}$  can change irregularly. Neutron removal is determined experimentally for each shield material as a combined removal coefficient. These are included in Table 4.1, for shields with a thickness of less than mean free paths and at least  $6 \text{ g/cm}^2$  of hydrogenous material, also included are values for shielding materials, only substance [28].

## II. ENVIROMENTAL IMPACT

A proton accelerator could impact on environment because of the prompt radiation and the emission of radioactive effluents, each of which may have an off-site radiological impact. A component of the direct radiation field is the *Skyshine*, due to the fact that the shielding in the vertical direction is not always constrained in this way so that more radiation (usually neutrons) may be emitted from the roof shielding of an accelerator at levels that may have an off-site impact.

Because the thresholds for nuclear reactions for neutrons with the constituents of air all lie near or above 20 MeV, the interactions below this energy are restricted to elastic scattering. The high energy nuclear interaction length for  $\text{N}_2$  and  $\text{O}_2$  are of the order of  $90 \text{ gcm}^{-2}$ , which for the density of air is of the order of 750 m and hence the high energy neutrons effectively escape to great distances. Because only low energy neutrons could be scattered backward, these neutrons predominate.

Due to mass ratio of neutrons to nitrogen and oxygen nuclei, many elastic scatters are needed in order to reduce the neutron energy. It follows that, as a first approximation, there is no effective attenuation of neutrons in air and the primary reduction in fluence

out to a few hundred meters derives from geometrical factors. Dependence of neutron dose on distance from the source is therefore in the first instance a purely geometrical effect. As particle number must be conserved, the dose is inversely proportional to the area over which the particles are dispersed

$$H = \frac{H_a A}{2\pi r^2} \quad (4.7)$$

where  $H_a \cdot A$  is the mean dose on  $A$  area of the roof and  $r$  is the distance from the source to the field point of interest [29].

### III. INDUCED RADIOACTIVITY

In an accelerator bunker for energies  $E < 30$  MeV, radionuclide production by direct reactions such as single and multi-nucleon transfer as well as processes such as  $(p, \gamma)$  are of principal concern. The systematic and approximate energy dependences of these processes are well understood. Reactions that occur inside bunker are endoergic nuclear reactions with a  $E_{th}$  equal to:

$$E_{th} = \frac{m_p + M}{M} |Q| \quad (4.8)$$

The Q-value is the difference between the separation energy of the in-going and out-going particles in the absence of excitation energy in either the entrance or the exit channels. It is also quite common for thermal neutrons to produce significant levels of induced radioactivity in the accelerator room. Such radioactivity results from thermal neutron capture reactions that sometimes can have relatively large cross sections. As the energy of the incident radiation increases, the number of possible reaction channels increases, with a corresponding increase in the number of radionuclides produced. The variety of radionuclides that can be produced becomes higher as one raises the bombarding energy because more reaction thresholds are exceeded. Induced radioactivity produced by neutron irradiation is more probable compared with gamma and charged particles. This is explained by the fact that gamma and charged particles must have high energy of at least a few MeV to start a nuclear reaction with the creation of radioactive nuclei, whereas neutrons are electrically neutral and easily penetrate the nuclei to make it radioactive. For protons or heavy ions, the nuclear rupture caused by the collision between the initial particle and the target material and the nuclear reaction between the generated secondary neutrons and the material are also important ways to generate induced radioactivity. Heavy ion projectiles are fragmented and remain implanted in the target after hitting it. Induced activity at modern proton and heavy ion accelerators is characterized by levels of the order of  $10^2 - 10^3 \mu\text{Sv/h}$  (depending on cooling times and irradiation scenarios) and causes significant challenges during commissioning and repair work. The ratio only applicable to induced radioactivity in air, water and unshielded accelerator structures, while the ratio in target or material test areas is often much higher. Sometimes, for such work, shielding is required to reduce radiation to levels that are acceptable in terms of dose limits and optimization of staff, or, with adequate consideration for the area's

intended level of human occupancy. The contribution from the induced activity of the concrete shielding to the total radiation background in the tunnels of the accelerator after it ceases can be significant, comparable to the background from the activation of various metal parts of the accelerator. The activation of soil and water ponds near accelerators creates significant problems. Due to their large dimensions and the level of prompt radiation generated during operation, high-energy accelerators are generally located underground. Following soil activation, generated radionuclides may leak and migrate into numerous streams and bodies of water. Neutrons passing through accelerator foundations constitute the most significant contribution to soil activation. The soil is often weakly activated, and its activation is determined by the long-lived radionuclides such as  $^3\text{H}$ ,  $^7\text{Be}$ ,  $^{22}\text{Na}$ ,  $^{45}\text{Ca}$ ,  $^{54}\text{Mn}$ ,  $^{55}\text{Fe}$ . The main contribution to the activation of ground water is made by  $^3\text{H}$  and  $^{22}\text{Na}$ , which form readily soluble compounds. In Table 4.2 are summarized radioactive nuclides produced in a particle accelerator.

**Table 4.1:** Neutron removal coefficients  $\Sigma_{nr}$  for thermal neutrons for materials which are surrounded by sufficient hydrogenous material to absorb neutrons that are degraded in energy due to scattering interactions.

Material	$\Sigma_{nr} (cm^{-1})$
Sodium	0.032
Graphite	0.078
Carbon	0.084
Concrete	0.089
Heavy Water	0.092
Zirconium	0.101
Water	0.103
Paraffin	0.106
Polyethylene	0.111
Lead	0.118
Beryllium	0.132
Iron	0.156
Copper	0.167
Uranium	0.182
Tungsten	0.212

**Table 4.2:** *Induced radioactivity in proton accelerators facilities [30].*

Material	Isotope	Threshold (MeV)	Half-life	$\sigma$ (mb)	Decay mode
Plastic, oils	$^3\text{H}$	11	12.33 y	10	$\beta^-$
-	$^7\text{Be}$	2	53.22 d	10	EC
Al	$^{22}\text{Na}$	30	2.60 y	10	$\beta^+$
Iron	$^{44m}\text{Sc}$	-	2.44 d	-	IT
-	$^{46}\text{Sc}$	-	83.8 d	-	$\beta^-$
-	$^{47}\text{Sc}$	-	3.35 d	-	$\beta^-$
-	$^{48}\text{Sc}$	-	1.82 d	-	$\beta^-$
-	$^{48}\text{V}$	20	15.97 d	6	$\beta^+$
-	$^{51}\text{Cr}$	30	27.7 d	6	EC
-	$^{52}\text{Mn}$	20	5.59 d	30	$\beta^+$
-	$^{54}\text{Mn}$	30	312.1 d	30	EC
-	$^{55}\text{Fe}$	-	2.74 y	-	EC
-	$^{59}\text{Fe}$	-	44.5 d	-	$\beta^-$
-	$^{56}\text{Co}$	5	77.2 d	30	$\beta^+$
-	$^{57}\text{Co}$	30	271.7 d	30	EC
-	$^{58}\text{Co}$	30	70.9 d	25	$\beta^+$
Steel	$^{59}\text{Ni}$	-	75 y	-	EC
-	$^{60}\text{Co}$	30	5.27 y	15	$\beta^-$
Copper	$^{63}\text{Ni}$	-	100 y	-	$\beta^-$
-	$^{65}\text{Zn}$	-	243.7 d	100	EC



## Chapter 5

# GEANT4 simulation

The GEANT4 (GEometry ANd Tracking) is a software coded in a object oriented programming language, C++ [31]. It can simulate the passage of particles through matter, with a lot of application in high energy, nuclear and accelerator physics, and medical and space science. It is made for detector design and response to particles, signal amplitudes, energy spectra, resolutions, efficiencies. The minimal tasks to provide in order to have a complete process are:

- provide geometrical and material description of apparatus
- provide event generator
- choose an appropriate physics list and production cuts

The software produce particles and simulate interactions, and propagate the secondary particles. At the heart of Geant4 is an abundant set of physics models to handle the interactions of particles with matter across a very wide energy range. Geant4 is written in C++ and exploits advanced software-engineering techniques and object-oriented technology to achieve transparency. In order to show the class description and utilization, is useful to put them in a diagram, as in figure 5.1. Here are showned categories and hereditary. So at the bottom of the diagram there are categories used by virtually all higher categories and provide the foundation of the toolkit.

**Global** category covers the system of units, constant, numeric and random number handling.

**Material and Particle** implement facilities necessary to describe the physical properties of particles and materials for the simulation of particle-matter interactions.

**Geometry** module offers the ability to describe a geometrical structure and propagate particles efficiently through it. Above these reside categories required to describe the tracking of particles and the physical processes they undergo.

**Track** category contains classes for tracks and steps, used by processes.



**Processes** category, which contains implementations of models of physical interactions: electromagnetic interactions of leptons, photons, hadrons and ions, and hadronic interactions.

**Tracking** category, which manages their contribution to the evolution of a track's state and provides information in sensitive volumes for hits and digitization.

**Event** category manages events in terms of their tracks.

**Run** category manages collections of events that share a common beam and detector implementation.

**Readout** category allows the handling of pile-up.

All of these categories use some capabilities to connect to facilities outside toolkit through abstract interfaces, in order to provide *visualization, persistency and user interface*.

## I. FIRST STEP SIMULATION

The aim of this simulation project is the study of ambient dose inside bunker, due to proton beam smashing on the matching line at the end of the RFQ-injector. So the first simulation is developed in order to obtain a contour map of the ambient dose produced by a pulse of the beam, and a mean number of secondary particles produced. In this starting phase, is designed not only the geometry, but also the electromagnetic field, in order to simulate the lost number of protons in the beam pipe for every tank. So at initial step isn't simulated the beam outside accelerator line, simulation is stopped at the end of Module 0.

### i. Geometry

The simulation of the accelerator LINAC of the ERHA project starts with the geometry construction of the matching line between the RFQ output and the input of *Module 0*, which specifications are listed in Table 5.1. The *Module 0* is designed in a simplified way,

**Table 5.1:** *Matching Line specifications.*

Element	Lenght	In. Radius	Ext. Radius	Material	Gradient
PMQ 0	20 mm	3 mm	10 mm	Alluminium	-160 T/m
Drift	65 mm	3 mm	3,2 mm	Stainless-Steel	-
PMQ 1	20 mm	3 mm	10 mm	Alluminium	196 T/m

in which tanks are cilindric and have same size for the entire module. This simplification is of course a limit, but as showned after field is resized and constant in the tank. So *Module 0* is descripted in Table 5.2. This assembly is copied 5 times and it is 1 meter long. The class in which is defined is *LXeAccelerator()*, written in Appendix A.

## ii. Materials

The beam pipe is the volume with a vacuum density material, and it is made with materials in Table 5.3 with a pressure of  $10^{-7}$  Pa. The classe in which materials are defined is *LXeDetectorConstruction()*, written in Appendix B.

## iii. Fields

The beam pipe is also the volume in which protons are accelerated, so here is defined the field. Because of difficulties in setting a non-uniform and non-constant field, the volumes of the beam-pipe inside the tanks are filled with electric field with a constant value along z-axis that is the mean value of the original time-dependent field in the tanks. Class *ElectricFieldSetup()* can perform this operation.

The beam pipe of the quadrupole is filled with a quadrupole magnetic field, thanks to *QuadrupoleMagneticField()* class used in *FieldSetup()*. In order to build a multithreading executable object, fields are defined *static* and *G4ThreadLocal*.

## iv. Results

A first interesting result is obtained by a *voxelitation* of the space of the envelope around accelerator. In this way is studied the dose released for a bunch of  $10^6$  protons. In figure 5.2 is clear that all the dose is released at the center in the beam pipe. In figure 5.3 dose is released at first and second quadrupole, as is showned also in figure 5.4.

## II. SECOND STEP SIMULATION

The second step of the simulation project is focused on a detector design. The gamma and neutron prompt radiation fields were measured by two different detectors. Simulation starts from the construction of detector, in order to simulate event detection of a gamma or a neutron.

**Table 5.2:** *Module 0 specifications.*

Element	Lenght	In. Radius	Ext. Radius	Material	Gradient(or $\vec{E}$ )
Tank 0	60 mm	1.5 mm	30 mm	Copper	4.8 MV/m
Drift	10 mm	3 mm	3.2 mm	Stainless-Steel	-
PMQ 2	20 mm	3 mm	10 mm	Alluminium	-179 T/m
Drift	10 mm	3 mm	3.2 mm	Stainless-Steel	-
Tank 1	60 mm	1.5 mm	30 mm	Copper	4.8 MV/m
Drift	10 mm	3 mm	3.2 mm	Stainless-Steel	-
PMQ 3	20 mm	3 mm	10 mm	Alluminium	186 T/m
Drift	10 mm	3 mm	3.2 mm	Stainless-Steel	-

i. Geometry

Detectors are placed near the accelerator at 1 meter, at same altitude of the beam-pipe as showned in figure 5.5. Simulation in this step is focused on the interaction between beam and materials around beam-pipe, and the tracks are stopped outside bunker without a interaction with the walls. Geometry description is different for gamma and neutron detectors: they are parallelepipedal shaped, gamma is  $6 \times 6 \times 20 \text{ cm}^3$ , while neutron is  $20 \times 20 \times 40 \text{ cm}^3$ . They consist of a scintillation material and a photomultiplier of  $2.3 \text{ cm}$  of radius on one side, covered by aluminium. Neutron detector is surrounded by  $3.0 \text{ cm}$  of polyethylene material. In figure 5.6 is visible the gamma detector.

ii. Materials

These dose monitor detectors are scintillating one (see Table 5.4). The glass scintillator inside neutron detector (density  $2.4 \text{ g/cm}^3$ ) is composed as described in Table 5.5.

iii. Results

First is simulated gamma detection, then neutron detection, by changing the *MainVolume* class.

**Gamma** Output is a table of simulated quantities (see Table 5.6), where *protons* is the number of particle injected, *hits* is the total number of optical photons that hit the pmt in the scintillator, *hits over threshold* is the total number of events detected, *scintillation ev*, is the number of optical photons produced by scintillation, *cherenkov ev.* is the number of optical photons produced by cherenkov effect, *absorbed ev.* is the number of optical photons absorbed by scintillator material, *absorbed energy* is the total energy absorbed in the scintillator and *integrated dose* is the total dose released in the simulated world.

In order to make a comparisation, linear extrapolation gives  $\sim 10^3$  gamma with respect to  $10^{11}$  protons, which is the number of protons accelerated in one second by injector. At this value correspond an equivalent dose of:

$$1 * 10^{-12} \text{ Sv} \cdot \text{cm}^2 * \frac{10^3}{\text{cm}^2 \cdot \text{s}} = 10^{-9} \frac{\text{Sv}}{\text{s}} = 3.6 \frac{\mu\text{Sv}}{\text{h}} \quad (5.1)$$

**Table 5.3:** Vacuum composition.

Element	Fraction
Oxygen	30%
Nitrogen	69%
Hydrogen	1%

**Neutron** The simulation of neutron detector is a little more complicated, because of the small rate of production. Indeed, in order to be sure is applied a window on total hits on pmt in scintillator detector, after a check on the peak position, like in figure 5.7. So when hits for simulated detector are  $800 < hits < 1200$ , it is a neutron. A first simulation is shown in Table 5.7, while a more statistically efficient simulation gives the results in Table 5.8. Again this value must be compared with the measurement. So with  $10^{11}$  protons per second, the number of neutrons could be obtained by a linear fit of data. Conversion to equivalent dose gives:

$$391 * 10^{-12} Sv \cdot cm^2 * \frac{170}{cm^2 \cdot s} = 66470 * 10^{-12} \frac{Sv}{s} = 239 \frac{\mu Sv}{h} \quad (5.2)$$

### III. THIRD STEP SIMULATION

The last step include the simulation of all the other modules up to 50 MeV of acceleration. The Module with CCL tanks is constructed with the use of a opensource library *CADMesh* [32]. For a simulation of the entire process is required a lot of CPU-time, as long as 5 days for  $10^9$  protons, with results shown in Table 5.9. The entire accelerator room is a bunker with thick walls (1.20 meters), of concrete material with a composition like in Table 5.10, as is shown in figure 5.8. A very important result is the study of energy of the beam for all the acceleration modules. In order to compute the energy of the beam is useful to plot the kinetic energy of the protons inside beam-pipe during a *step*, a GEANT4 object that memorize several parameteres between two interaction. In figure 5.9 is shown in the x-axis the energy, while on the y-axis there is number of steps.  $N(T)$  is the number of steps at  $T$  kinetic energy, and this is proportional to  $N_p(T)$  that is the number of protons, proportional to  $I(T)$ , current of the beam:

$$\frac{N(T)}{N(0)} = \frac{N_p(T)}{N_p(0)} = \frac{I(T)}{I(0)} \quad (5.3)$$

where  $I(0)$  is the mean peak current of the RFQ. At the end of *Module 0* the current is:

$$I(T) = 0.25 * 1.2mA = 300\mu A \quad (5.4)$$

As shown results aren't statistically good, because of CPU-time consumption. So in order to reduce time is possible to change run from *singlethread* to *multithread*.

**Table 5.4:** *Scintillator specifications: R.Ind is the refractive index, Abs. Lenght is the absorbtion lenght, Yield is the number of gamma per unit of energy.*

Scintill.	R.Ind.	Abs. Lenght	Yield	$\tau$	Birks constant
Polystyrene	1.58	250 cm	10 g/keV	2.4 ns	0.126 mm/MeV
Glass	1.59	420 cm	10 g/keV	18 -/- 45 ns	0.126 mm/MeV

### i. Multithreading

The multithreading approach is obtained using *G4MTRunManager* class. Use of shared object defined *static* and *G4ThreadLocal* in *LXeDetectorConstruction.hh*, gives rise to a memory reduction, but in primis delete errors during run due to field and geometry. It needs 12 hours for a run with  $10^9$  protons, a very time-consumption reduction. In Table 5.11 are shown results of simulation and in Figure 5.10 is shown the linear fit used to derive the coefficient for the expected dose rate.

### ii. Superposition of sources

As a consequence of the high number of protons accelerated by simulation with respect to real current at the end of module 7, as is shown in figure 5.9, a new method of simulation could be a separation between injection and beam output in air. This could save a lot of time, because very high time-consumption is needed for simulation in air.

**Table 5.5:** *Glass scintillator specifications.*

Element	Fraction
C	84.18%
H	7.82%
Li6	7.5%
Ce	0.5%

**Table 5.6:** *Results with gamma detector.*

Protons	$10^9$
Hits	1250
Hits over threshold	11
Scintillation ev.	3100
Cherenkov ev.	0
Absorbed ev.	1870
Absorbed Energy	1170 keV
Integrated dose	$1.1 * 10^{-12}$ Gy

**Table 5.7:** *Second step neutron simulation  $10^9$  protons.*

Protons	$10^9$
Hits	17600
Hits over threshold	17
Scintillation ev.	38600
Cherenkov ev.	704
Absorbed ev.	21400
Absorbed Energy	9357 keV
Integrated dose	$1.02 * 10^{-12}$ Gy
Neutrons	1

**Table 5.8:** *Second step neutron simulation  $10^{10}$  protons.*

Protons	$10^{10}$
Hits	3729000
Hits over threshold	2200
Scintillation ev.	8206000
Cherenkov ev.	70000
Absorbed ev.	4500000
Absorbed Energy	1243000 keV
Integrated dose	$1.2 * 10^{-11}$ Gy
Neutrons	17

**Table 5.9:** *Third step neutron simulation.*

Protons	$10^9$
Hits over threshold	34
Neutrons	3

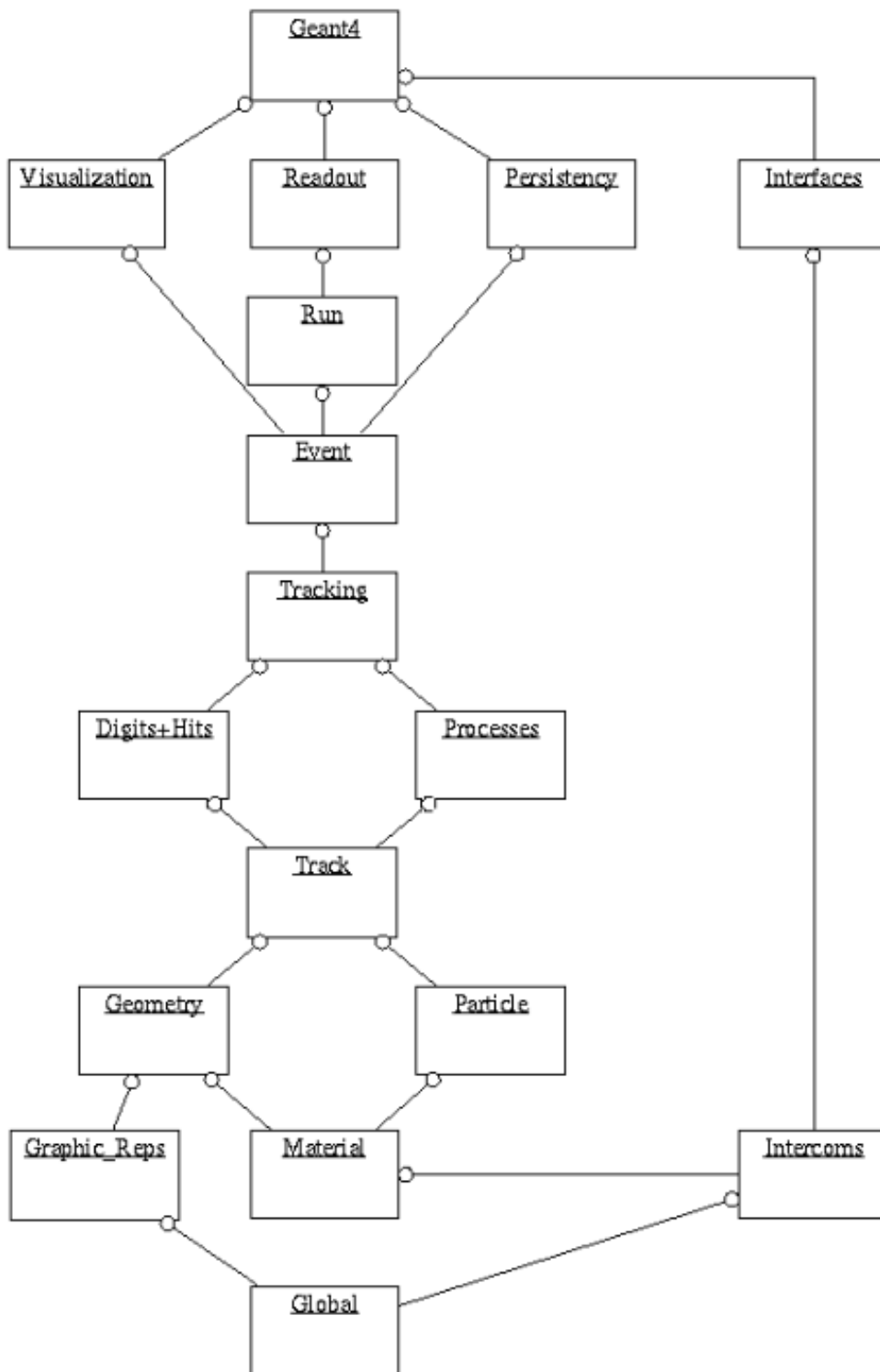


Figure 5.1: Geant4 class category diagram.

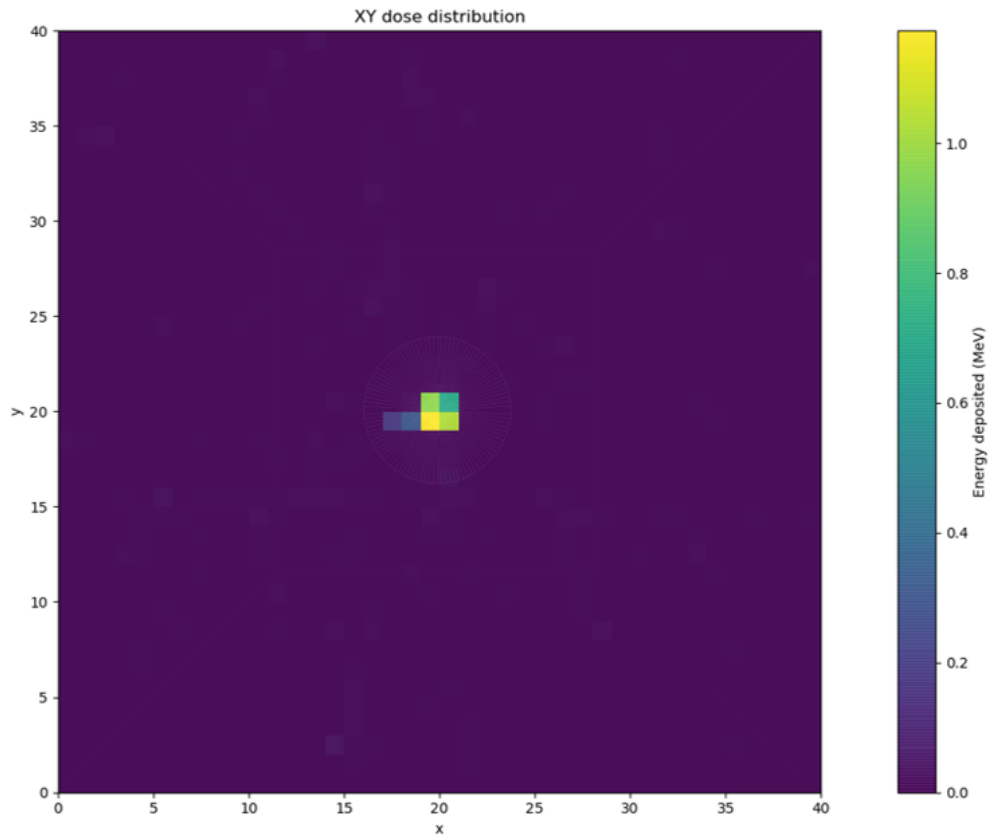


Figure 5.2: The map of the dose in  $xy$  plane.

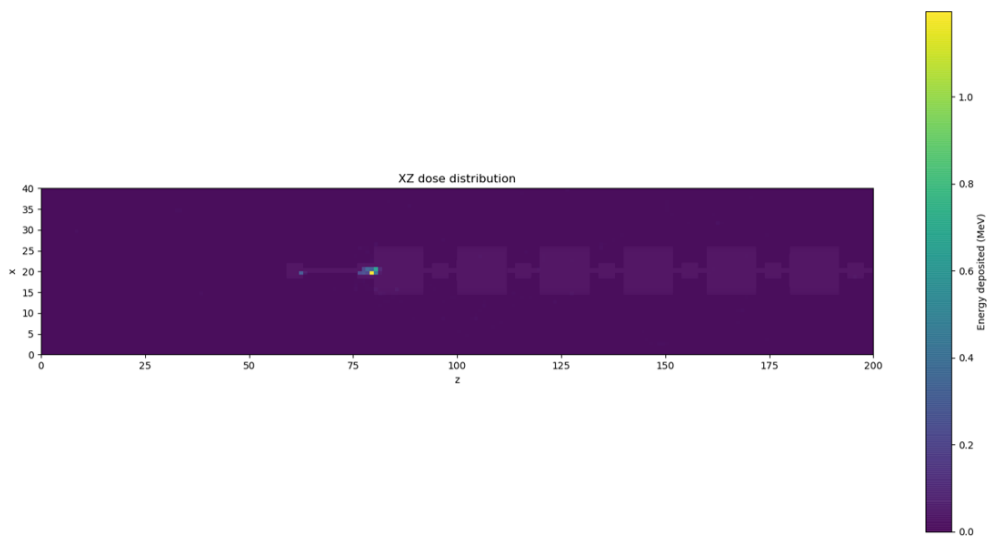


Figure 5.3: The map of the dose in  $xz$  plane.



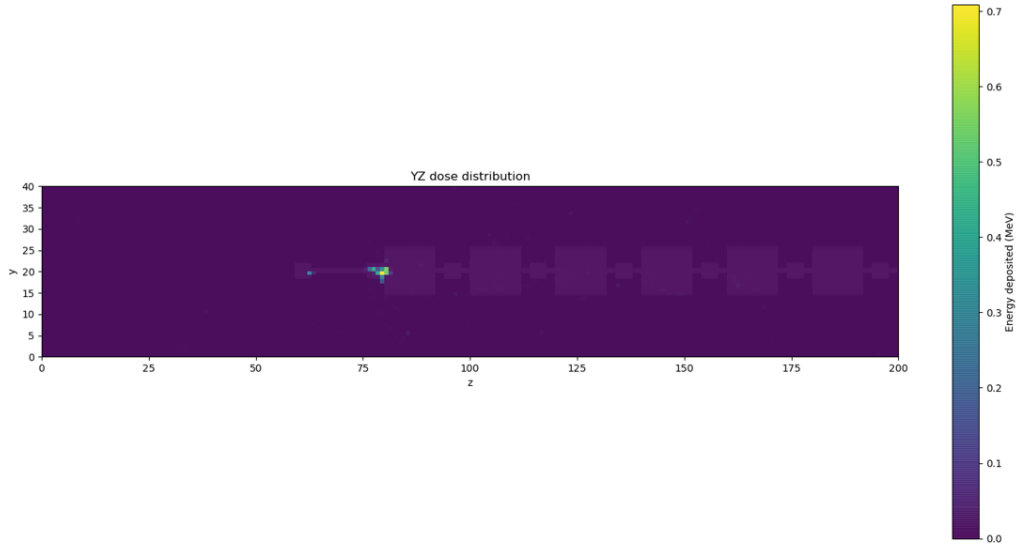


Figure 5.4: *The map of the dose in yz plane.*

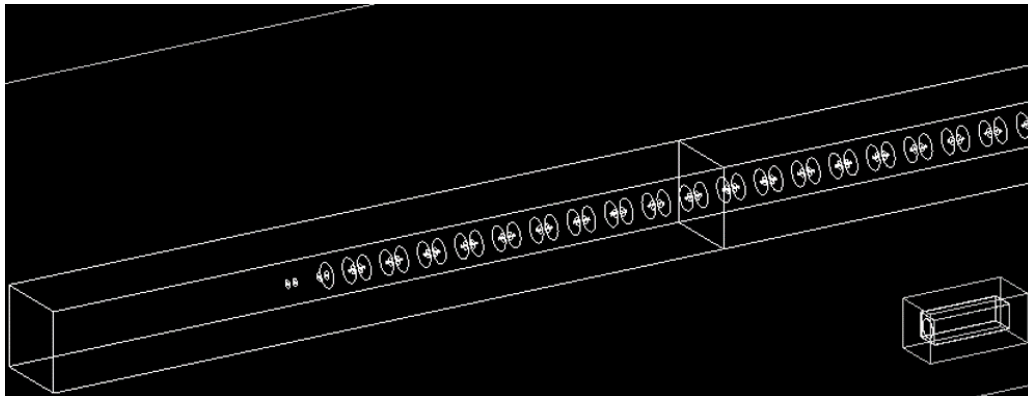
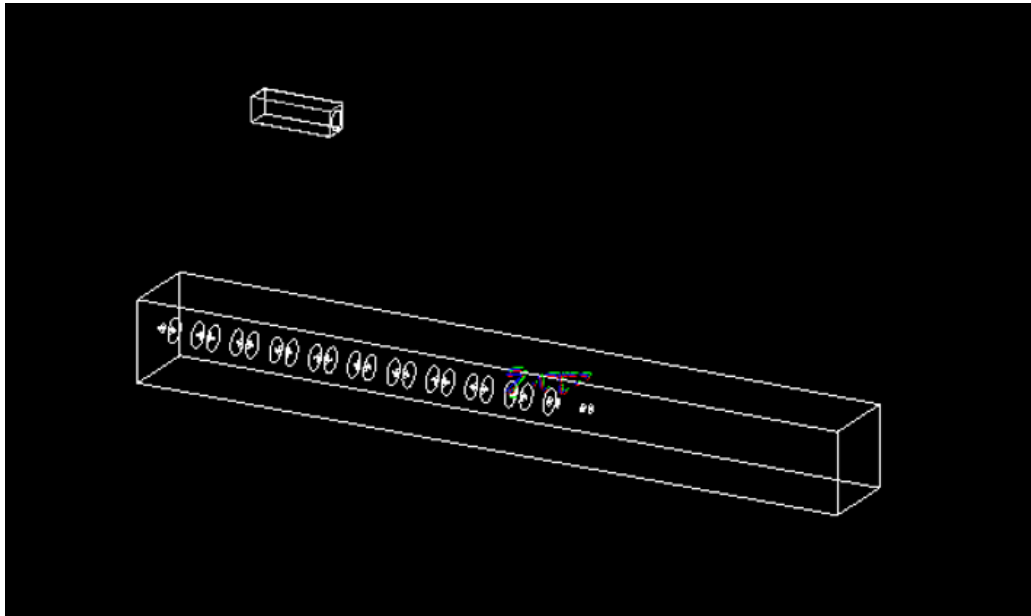


Figure 5.5: *A screen of the neutron detector position.*



**Figure 5.6:** A screen of the gamma detector position.

**Table 5.10:** Concrete composition.

Oxygen	49.2875%
Carbon	5.62%
Hydrogen	0.6%
Sodium	0.453%
Magnesium	0.663%
Alluminium	2.063%
Silicon	18.867%
Potassium	0.656%
Calcium	20.091%
Iron	1.118%
Phosphorus	0.048%
Sulphur	0.012%
Titanium	0.347%
Manganese	0.0387%
Zinc	0.0241%
Zirconium	0.0074%
Barium	0.0179%
Lead	0.0464%
Strontium	0.04%

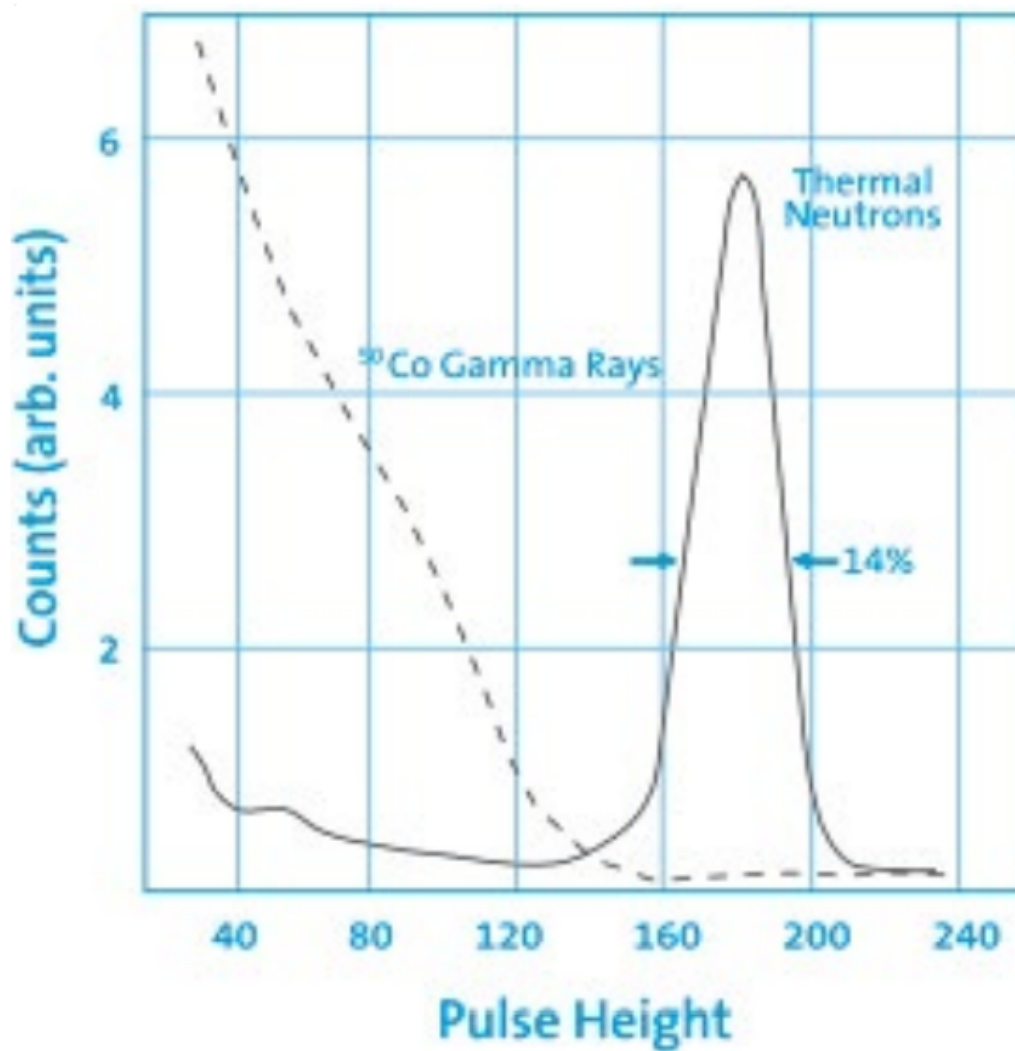
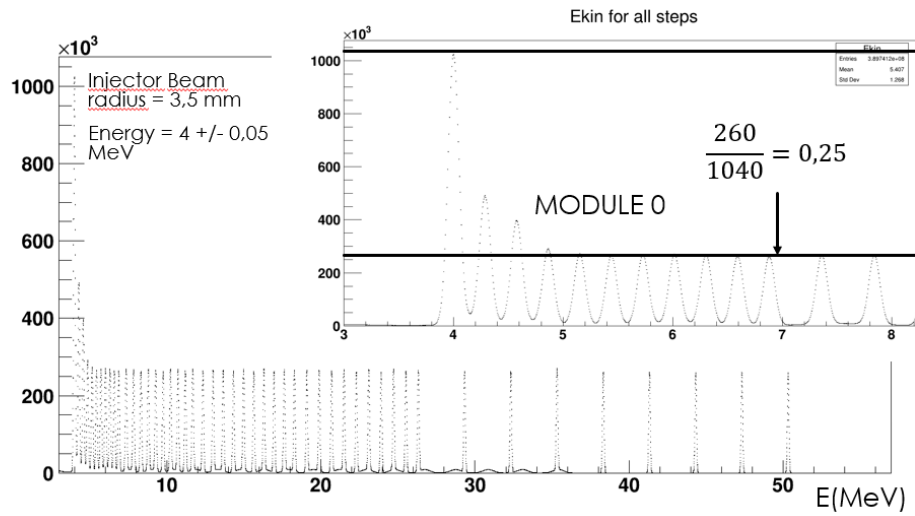


Figure 5.7: Pulse height curve of a glass scintillator filled with lithium.



Figure 5.8: Top to bottom view of the simulated bunker.

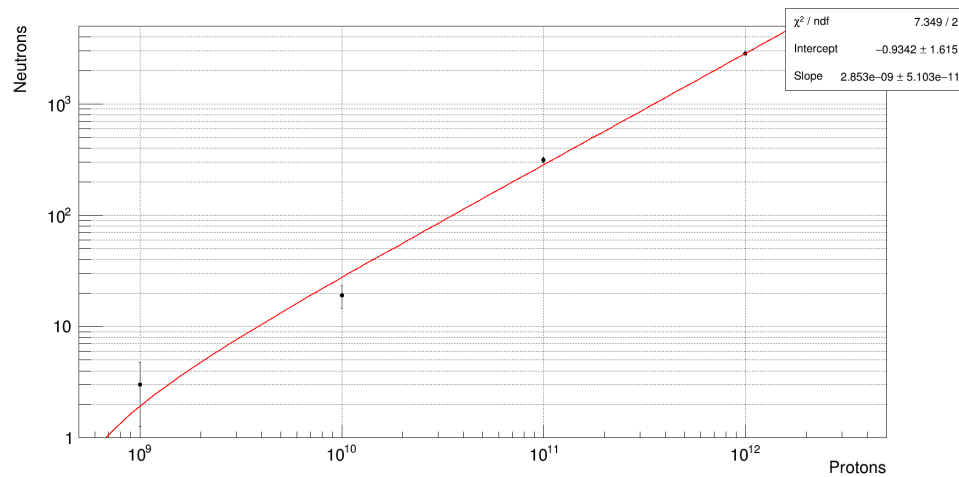
### Section III



**Figure 5.9:** The number of steps inside beam-pipe related to kinetic energy of the beam.

**Table 5.11:** Number of neutrons into detector simulated for different numbers of protons.

Protons	Neutrons
$10^9$	3
$10^{10}$	19
$10^{11}$	315
$10^{12}$	2836
Intercept	$(-0,9342 \pm 1,6150) n$
Slope	$(2,853 \cdot 10^{-9} \pm 5,103 \cdot 10^{-11}) n/p$



**Figure 5.10:** Linear fit, parameter extraction with MIGRAD-ROOT.

## Section III

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## Chapter 6

# Gamma and Neutron detection

Detection of particles and dose measurement in a pulsed neutron field is a issue for particle accelerators, where beam is lost near targets, collimators and beam dumps. The interest in active detectors to be employed in pulsed neutron field is constantly increasing due to the growing number of applications where the pulse structure of the radiation field hinders the use of active detectors operating in pulse mode in the halls housing acceleratos, including the treatment rooms.

Although the  $H^*(10)$  which characterises a single burst is usually low, and would no constitute a problem in terms of averaged  $H^*(10)$  rate over the entire measurement period, the  $H^*(10)$  rate during the radiation burst can reach extremely high values, up to 100 Sv/h in typical medical diagnostic applications and up to  $10^7$  Sv/h [33] in facilities such as the ones used for material testing, and this usually leads to severe underestimations of the  $H^*(10)$ . Recent measurements around research particle accelerators reported severe under reponse in commercial active neutron detectors.

Main cause of this underestimation can be attributed to dead time losses, which are a distinctive feature of active detectors working in pulse mode. An active radiation detector can in fact operate in three modes: current, mean square voltage (MSV) and pulse mode. Current mode averages out the fluctuations in the intervals between individual interactions and is usually employed with high interaction rates when there is no need of preserving the information on the amplitude and timing of single interactions. MVS mode, whose detection principle is based on a special processing of the fluctuating component of the detector current signal, becomes useful when making measurements in mixed radiation environments when the charge produced by one type of radiation is much different than that from the second type. Pulse mode is the most commonly applied, for preserving information on the amplitude and timing of individual events. In this detector each pulse represents the results of a single interaction. Disadvantage of this property is dead time and losses induced by the counting system, which can be correctly compensated only in the case of steady-state sources of constant intensity, but not in the case of pulsed sources. An ideal detector would in fact count every event that occurs. However, a real detector and its read-out electronics need a specific amount of time to create and process an output pulse. An event that occurs during this time cannot be

registered. Depending on the detector system, it is either suppressed (non paralyzable systems) or changes the shape of the previously detected pulse, resulting in a pile up (paralyzable systems)[34]. The minimal time between two separately detectable events is called the dead time and it ranges from 1 to 10  $\mu\text{s}$  for neutron dose detectors. It is possible to define five requirements that an active neutron detector should show for efficiently working in pulsed neutron field:

- capability to withstand very high instantaneous neutron fluxes with little or no saturation;
- high sensitivity, usually expressed in  $\text{nSv}^{-1}$ , at least comparable with that of commercially available rem counters, i.e. about  $1 \text{ nSv}^{-1}$ ;
- capability to reject the photon contribution that usually accompanies the neutron fields;
- capability to measure correctly the intensity of a single neutron burst;
- good sensitivity over the entire neutron energy range, especially to high-energy neutrons.

### I. DEAD-TIME CORRECTION FACTOR

The difficulty which arises in electrical counting when the source is pulsed is due to the finite resolution of any counting system. Thus, following any count, there is a dead-time during which the system would be unable to record a further count, should one occur. It needs a careful study of the counting losses expected under that situation. It is seen that one cannot simply apply equations for continuous counting at the enhanced counting speed, the error in doing so would be greatest for conditions which are very likely to arise in practice, for example, when the pulse length does not greatly exceed the dead-time of the system. It is also shown that it is desirable to reduce the dead-time to the minimum. The studies of C. H. Westcott ([35]) on the suppression factor introduced when  $\frac{T}{\tau}$  ( $T$  is the pulse period), although greater than unity, cannot be considered large, lead to the simple formula

$$E' = E \frac{1}{X} \left( \frac{1}{2} + \frac{X - \frac{1}{2}}{1 + z} \right) \quad (6.1)$$

where:

- $E$  is the expected total count, including suppressed counts;
- $E'$  is the expected total of recorded counts;
- $X$  is  $\frac{T}{\tau}$  the duration of the pulse in units of the dead-time;
- $z$  is  $\nu\tau = \frac{N}{T}\tau$  is the expected mean counting rate during the pulse ( $\nu$ ) in units of the dead-time, indeed the average expectation of counts within one dead-time.

The factor for the standard condition of 0,6 mA of injector current could be calculated for a dead-time value of 1,0  $\mu\text{s}$  as measured in [36], and it gives

$$\frac{E'}{E} = \frac{1\mu\text{s}}{4\mu\text{s}} \left( \frac{1}{2} + \frac{4 - \frac{1}{2}}{1 + \frac{43}{4\mu\text{s}} 1\mu\text{s}} \right) \sim 0,2 \quad (6.2)$$

while for a current of 1,2 mA it gives 0,17.

## II. MEASURED EQUIVALENT DOSE

of accelerator started with an evaluation of gamma dose recorded in real time with a safety system monitor builded by *Atomtek*, a plastic scintillator detector. In order to measure the neutron dose in the bunker are set 3 different neutron probe in different positions. *Atomtek* built the first detector, a proportional counter with  $^3\text{He}$  in a polyethylene moderator, which is set using an FPGA with a RS232 protocol port by which are inserted commands using a LabView interface. After a first measure, a complete set of measurement is obtained with a *Thermo-BIOREM 752* probe, a proportional counter with  $\text{BF}_3$  placed in a cylindrical moderator containing polyethylene and boron carbide.

### i. Gamma Atomtek BDKG-04

*Atomtek BDKG-04* x-ray and gamma probe measures equivalent ambient dose and equivalent dose rate with features specified in Table 6.1 and shown in Figure 6.1. The monitor

**Table 6.1:** *Atomtek BDKG-04 features.*

Detector	scintillation plastic
Dose equivalent rate	0,05 $\mu\text{Sv/h}$ - 10 Sv/h
Intrinsic measurement error	$\leq 20\%$
Energy range	15 keV - 3 MeV
Energy sensitivity response	15 - 60 keV $\pm 35\%$ 60 keV - 3 MeV $\pm 20\%$
Sensitivity on $^{137}\text{Cs}$	70 cps for 1 $\mu\text{Sv/h}$
Operating temperature range	$-30^\circ\text{C}$ + $50^\circ\text{C}$
Relative humidity	$35^\circ\text{C}$ up to 98%
Protection class	IP64
Radio disturbance	CEI/IEC CESP 22:1997
Electromagnetic compatibility	CEI/IEC 61000-4-2:1995 IEC 61000-4-3:1995
Weight	0.45 kg
Dimensions	$\varnothing$ 60x200 mm

system consists of a set of gamma-probes located in different positions in the bunker. In Figure 6.2 is shown where are placed the detectors. The detector is placed at 1 meter from



beam-pipe at link point between Module 0 and Module 1. Gamma measurements are summarized in Table 6.2.

**Table 6.2:** Measured gamma dose for  $10^{11}$  protons.

Protons	$10^{11}$
Dose rate	$3,07\mu\text{Sv/h}$

## ii. Neutron Atomtek BDKN-03

Neutron detector BDKN-03 is a proportional counter with  $^3\text{He}$  in a polyethylene moderator, as depicted in Table 6.3 where are summarized the detector features. In Figure 6.3 is

**Table 6.3:** Atomtek BDKN-03 features.

Detector	$^3\text{He}$ proportional counter
Dose equivalent rate range	$0,1\mu\text{Sv/h} - 10 \text{ mSv/h}$
Dose equivalent range	$0,1\mu\text{Sv} - 10 \text{ Sv}$
Intrinsic measurement error	$\pm 20\%$
Energy range	$0,025 \text{ eV} - 14 \text{ MeV}$
Sensitivity to Pu–Be (dose mode)	$0.355 \text{ cps for } 1\mu\text{Sv/h}$
Flux density range	$0.1 - 10^4 \text{ neutron}\cdot\text{s}^{-1} \cdot \text{cm}^{-2}$
Sensitivity to Pu–Be (flux mode)	$0.5 \text{ cps for } 1\mu\text{Sv/h}$
Operating temperature range	$-30^\circ\text{C} + 50^\circ\text{C}$
Relative humidity	$\leq 35^\circ\text{C up to } 95\%$
Protection class	IP64
Weight	8 kg
Dimensions	$316\times 220\times 265 \text{ mm}$

shown a picture of the detector with LabView interface. Measured dose rate is plotted in Figure 6.4, 6.5 and 6.6. In Table 6.4 are summarized measured dose rate parameters. The analysis is made with RooFit, with a Gaussian Model for the signal and a Uniform model for the background.

## iii. Neutron ThermoScientific-BIOREM 752

The ThermoScientific FHT 752 BIOREM in Figure 6.7 is a commercial neutron dose rate meter for stationary and portable use, especially suited for environmental measurements.

In Table 6.5 the probe features are summarized. It employs a  $\text{BF}_3$  proportional counter placed in a cylindrical moderator containing polyethylene and boron carbide. The output is given in  $H^*(10)$ , but an internal calibration factor, expressed in  $\text{nSv/count}$ , can be set by the user, by calibrating it with a neutron source. The response function is given in Figure 6.8.

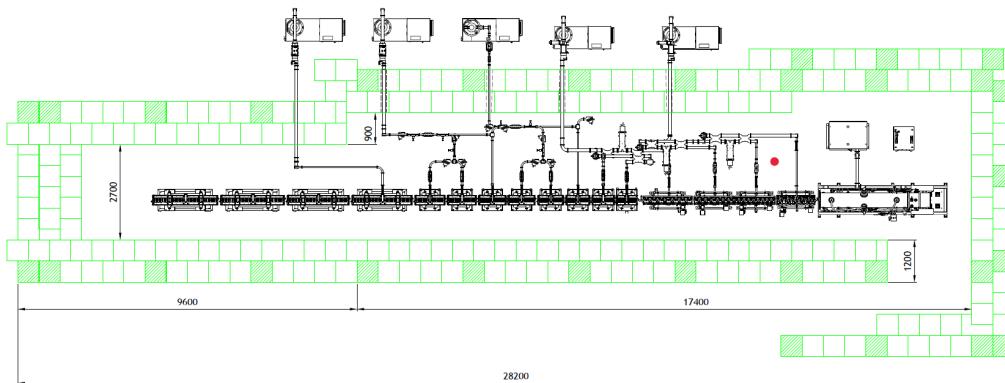
**Linearity plot** The expected  $H^*(10)$  is calculated by applying to the beam intensity a coefficient, in  $\text{n/p}$ , neutron per proton, derived from a linear fit of the simulated neutron detection in Figure 5.10, and then by calculating the simulated dose with equation 5.2, and in the end by applying the dead-time factor in equation 6.1. In Table 6.6 are shown these values for every beam intensities. The results of the measurements are shown in Table 6.7. In Figure 6.9 is shown the linearity plot[36]. The lines fitting the experimental data points are compared with a straight line that represents the bisector of the first quadrant, i.e. the ideal linear response. The equation used to fit the data were chosen only as visual guides and are linear of the form  $y = A \cdot x$ . The measured values refer to the dose rate  $H^*(10)$  detected in the stray field generated by a sequence of single beam pulses with a fixed repetition frequency. Uncertainty is statistical.

**Table 6.4:** Measured dose parameters: second value is more precise than first one because it is obtained in floating mean mode.

Current	Frequency	Dose rate	Standard Deviation
0.6 mA	10 Hz	133,67 $\mu\text{Sv/h}$	36,64 $\mu\text{Sv/h}$
0.6 mA	10 Hz	134,66 $\mu\text{Sv/h}$	7,01 $\mu\text{Sv/h}$
1.0 mA	10 Hz	200.05 $\mu\text{Sv/h}$	15,01 $\mu\text{Sv/h}$
1.2 mA	10 Hz	224,49 $\mu\text{Sv/h}$	51,52 $\mu\text{Sv/h}$



**Figure 6.1:** *X-ray and gamma probe.*



**Figure 6.2:** *Model shows the position of gamma and neutron detectors (red point), at 1 meter from beam-pipe and between Module 0 and Module 1.*

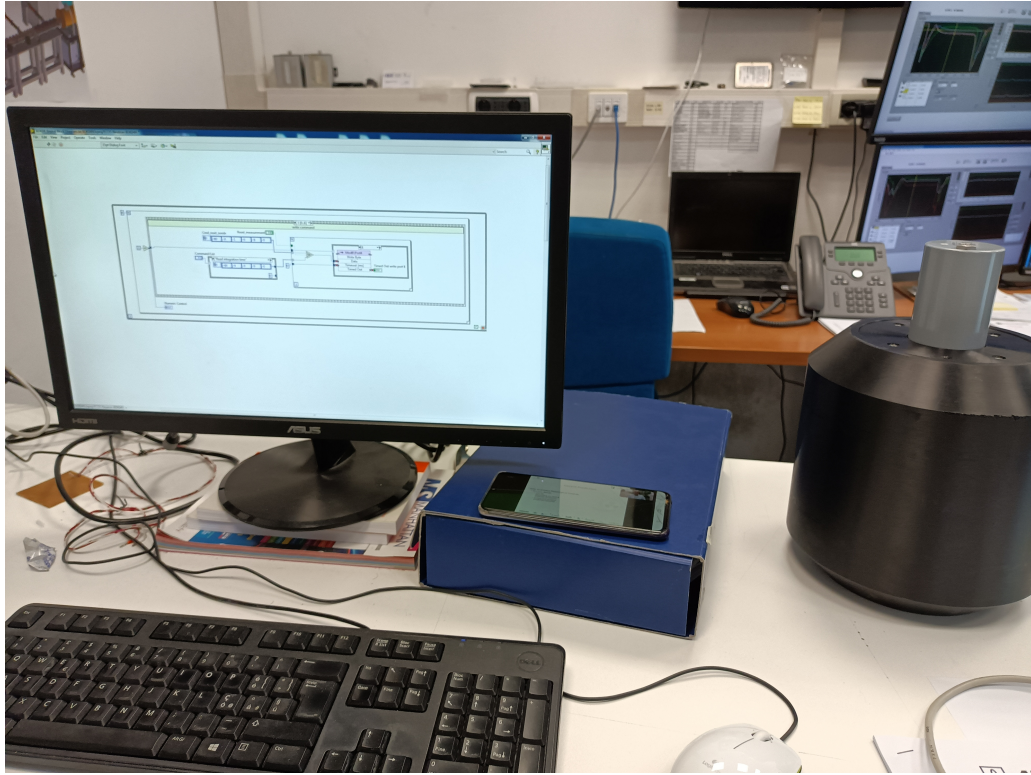


Figure 6.3: Atomtek neutron probe and Labview interface with RS232 protocol.

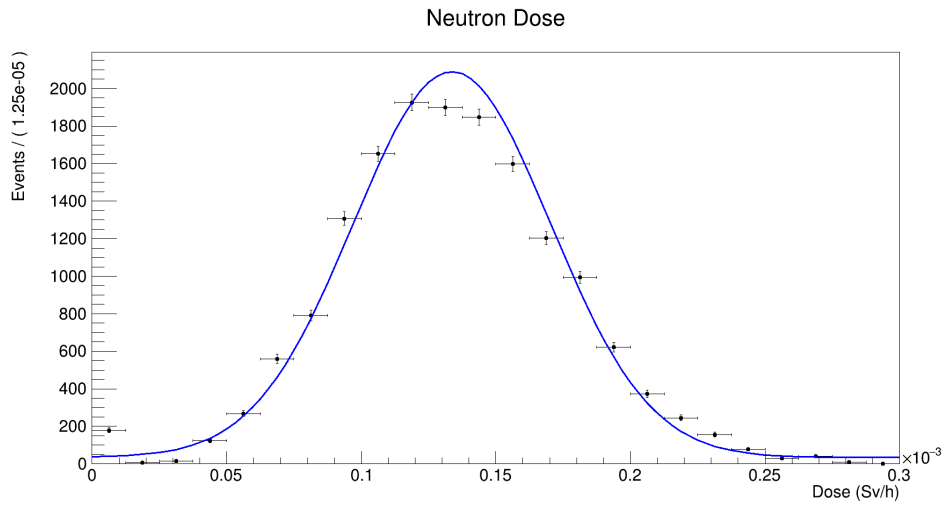
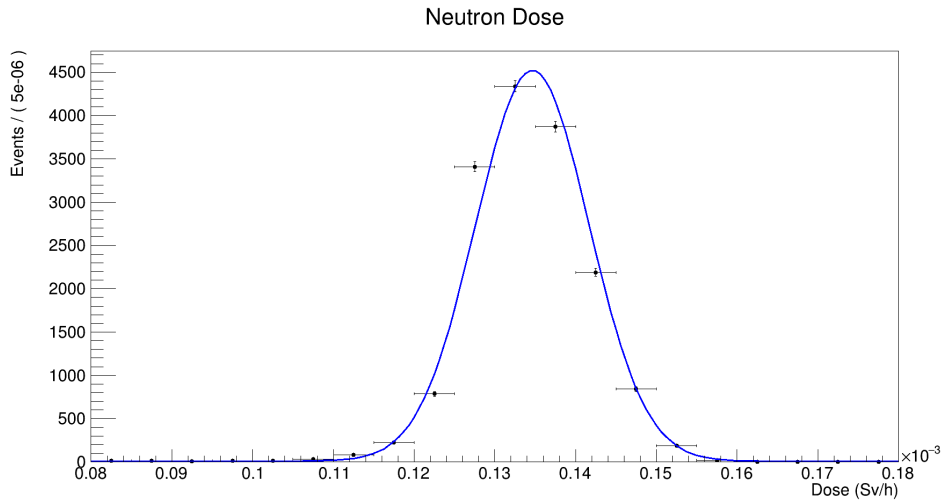
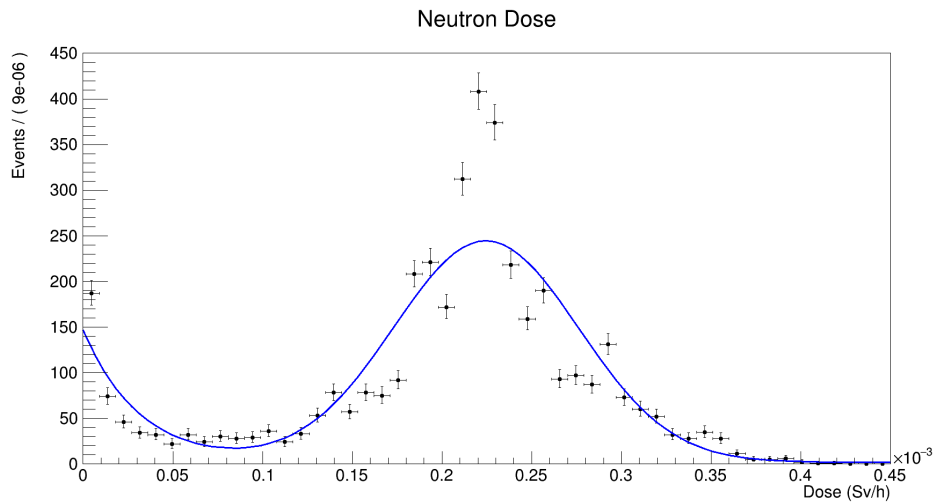


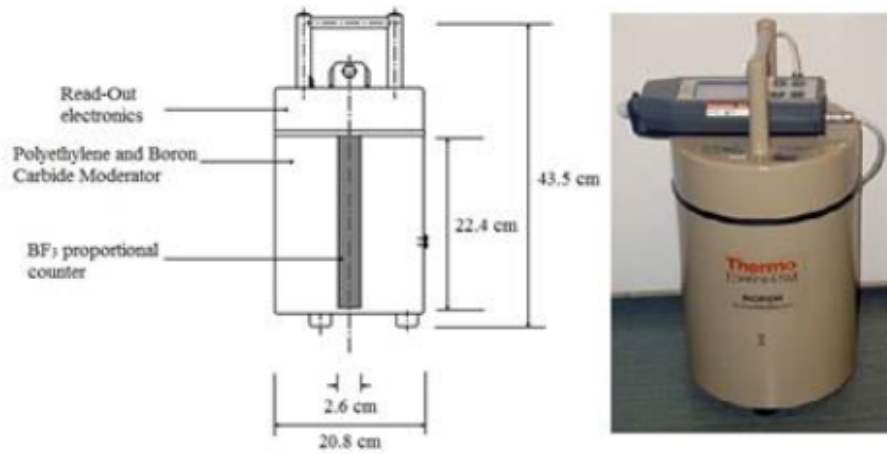
Figure 6.4: Neutron dose rate for an injection current of 0.6 mA and 10 Hz of repetition rate.



**Figure 6.5:** Neutron dose rate for an injection current of 0.6 mA and 10 Hz of repetition rate, in floating mean acquisition mode.



**Figure 6.6:** Neutron dose rate for an injection current of 1.2 mA and 10 Hz of repetition rate. Fit model is a Gaussian for signal, a Uniform for background and a Exponential for low rate events.



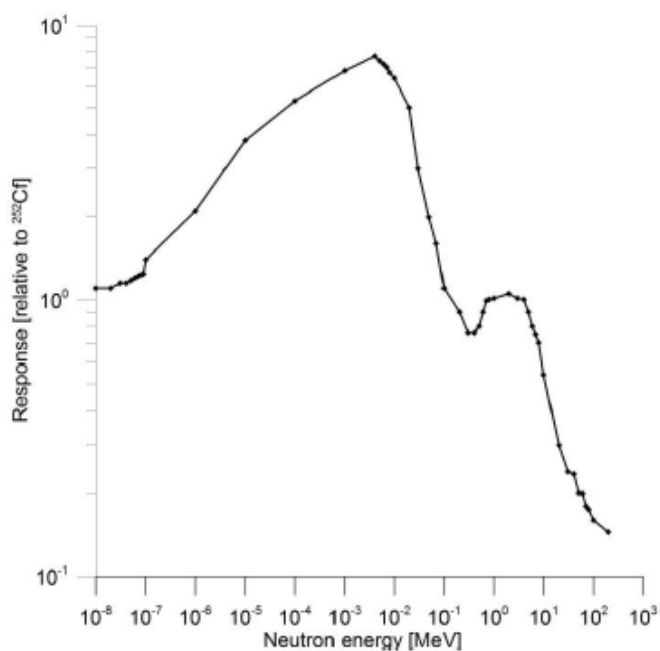
**Figure 6.7:** Geometry structure to the left, a picture to the right.

**Table 6.5:** Thermo-Scientific BIOREM FHT 752.

Detector	BF <sub>3</sub> proportional counter
Dose equivalent rate range	0,0001 $\mu$ Sv/h - 400 mSv/h
Intrinsic measurement error	$\pm$ 20%
Energy range	0,025 eV - 10 MeV
Sensitivity to <sup>252</sup> Cf (dose mode)	0.56 cps for 1 $\mu$ Sv/h
Protection class	IP64
Weight	11,5 kg
Dimensions	$\varnothing$ 208x435 mm

**Table 6.6:** Calculated expected dose rate values from the simulated ones.

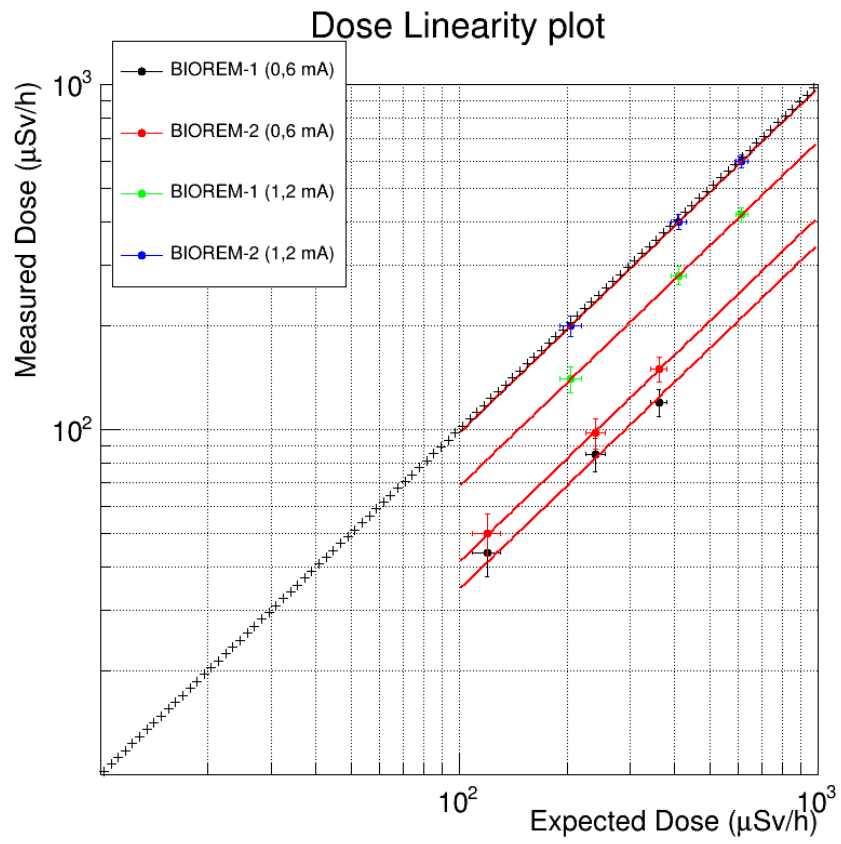
I (mA)	$\nu$ (Hz)	P/pulse	P/s	N/s	S. Dose ( $\mu$ Sv/h)	E. Dose ( $\mu$ Sv/h)
0.6	10	1,5 10 <sup>10</sup>	1,5 10 <sup>11</sup>	428	602,45	120,5
0.6	20	1,5 10 <sup>10</sup>	3 10 <sup>11</sup>	856	1204,90	241,0
0.6	30	1,5 10 <sup>10</sup>	4,5 10 <sup>11</sup>	1284	1807,35	361,5
1.2	10	3 10 <sup>10</sup>	3 10 <sup>11</sup>	856	1204,9	205,0
1.2	20	3 10 <sup>10</sup>	6 10 <sup>11</sup>	1712	2409,8	409,7
1.2	30	3 10 <sup>10</sup>	9 10 <sup>11</sup>	2568	3614,7	614,5



**Figure 6.8:** Response function to neutrons of BIOREM, expressed in units relative to the response of moderated <sup>252</sup>Cf [34].

**Table 6.7:** Measured dose rate values from two Thermo BIOREM probes.

I (mA)	$\nu$ (Hz)	BIOREM-1 ( $\mu$ Sv/h)	BIOREM-2 ( $\mu$ Sv/h)
0.6	10	44	50
0.6	20	85	98
0.6	30	120	150
1.2	10	140	200
1.2	20	280	400
1.2	30	420	600



**Figure 6.9:** Linearity plot for the BIOREM-1 and BIOREM-2 probes.





# Conclusion and Outlook

In order to resume this work, it is fundamental to say that simulation plays a central role in every situation in which there are radiation fields propagating where people have to be kept in safe. The reason is that particles such as neutrons behave in a way that is difficult to roughly estimate, and are responsible for the main part of the amount of dose absorbed by tissue. Shieldings are required to protect body and materials from damage, but they are very useful where there are steady-state field, so a radiation coming from nuclear decay. But in an accelerator facility there are other sources of neutrons and gamma, in general pulsed radiation fields, which are difficult to estimate. This work is focused to obtain a estimation of exposure dose due to the interaction between protons and materials around beam-pipe. Simulation is obtained with different approaches and than is compared with measured values of dose obtained with different probes. Results are very similar, if a factor that is used to get rid of undetected particles is applied, because of detector dead-time. Finally this work is useful to establish work conditions and verify measurements with probes. In order to go on with this project it could be useful to improve the simulation with a multithreading process and run it on a cluster for HPC. A further study on the proton beam and its interaction in air is of fundamental importance for an overall study of the dose released by the machine.



## Appendix A

# LXeAccelerator

GEANT4, C++ LXeAccelerator.cc class:

```
1 //
2 // *****
3 // * License and Disclaimer *
4 // *
5 // * The Geant4 software is copyright of the Copyright Holders of *
6 // * the Geant4 Collaboration. It is provided under the terms and *
7 // * conditions of the Geant4 Software License, included in the file *
8 // * LICENSE and available at http://cern.ch/geant4/license . These *
9 // * include a list of copyright holders. *
10 // *
11 // * Neither the authors of this software system, nor their employing *
12 // * institutes, nor the agencies providing financial support for this *
13 // * work make any representation or warranty, express or implied, *
14 // * regarding this software system or assume any liability for its *
15 // * use. Please see the license in the file LICENSE and URL above *
16 // * for the full disclaimer and the limitation of liability. *
17 // *
18 // * This code implementation is the result of the scientific and *
19 // * technical work of the GEANT4 collaboration. *
20 // * By using, copying, modifying or distributing the software (or *
21 // * any work based on the software) you agree to acknowledge its *
22 // * use in resulting scientific publications, and indicate your *
23 // * acceptance of all terms of the Geant4 Software license. *
24 // *****
25 //
26 //
27 /// \file optical/LXe/src/LXeMainVolume.cc
28 /// \brief Implementation of the LXeMainVolume class
29 //
30 //
31 #include "LXeAccelerator.hh"
32
33 #include "G4Box.hh"
34 #include "G4Colour.hh"
35 #include "G4LogicalSkinSurface.hh"
```

```
36 #include "G4LogicalBorderSurface.hh"
37 #include "G4LogicalVolume.hh"
38 #include "G4Material.hh"
39 #include "G4MaterialPropertiesTable.hh"
40 #include "G4OpticalSurface.hh"
41 #include "G4Sphere.hh"
42 #include "G4SystemOfUnits.hh"
43 #include "G4Tubs.hh"
44 #include "G4VisAttributes.hh"
45 #include "ElectricFieldSetup.hh"
46 #include "F03FieldSetup.hh"
47 #include "G4TransportationManager.hh"
48 #include "G4UniformMagField.hh"
49 #include "G4FieldManager.hh"
50 #include "G4RunManager.hh"
51 #include "G4NistManager.hh"
52 #include "G4Cons.hh"
53 #include "G4Orb.hh"
54 #include "G4Trd.hh"
55 #include "G4PVPlacement.hh"
56 #include "G4AutoDelete.hh"
57 #include "G4AssemblyVolume.hh"
58 #include "G4Types.hh"
59
60 //.....oooOOOOooo.....oooOOOOooo.....oooOOOOooo.....oooOOOOooo
61 .....
62 LXeAccelerator::LXeAccelerator(G4RotationMatrix* pRot, const G4ThreeVector&
    tlate ,
63 G4LogicalVolume* pMotherLogical, G4bool pMany,
64 G4int pCopyNo, LXeDetectorConstruction* c)
65 // Pass info to the G4PVPlacement constructor
66 : G4PVPlacement(pRot, tlate ,
67 // Temp logical volume must be created here
68 new G4LogicalVolume(new G4Box("temp2", 1.*cm, 1.*cm, 1.*cm),
69 G4Material::GetMaterial("Vacuum"), "temp2",
70 0, 0, 0),
71 "housing2", pMotherLogical, pMany, pCopyNo)
72 , fConstructor(c)
73 {
74 // Get nist material manager
75 G4NistManager* nist = G4NistManager::Instance();
76
77 // Envelope parameters
78 //
79 G4double env_sizeXY = 20.*cm, env_sizeZ = (0.99+0.80)*m;
80 G4Material* env_mat = G4Material::GetMaterial("Air");
81
82 // Option to switch on/off checking of volumes overlaps
83 //
84 G4bool checkOverlaps = true;
85
```

```
86 //
87 // Envelope
88 //
89 G4Box* solidEnv =
90 new G4Box("Envelope", //its name
91 0.5*env_sizeXY, 0.5*env_sizeXY, 0.5*env_sizeZ); //its size
92
93 G4LogicalVolume* logicEnv =
94 new G4LogicalVolume(solidEnv, //its solid
95 env_mat, //its material
96 "Envelope"); //its name
97
98
99 //Material - Air vacuum
100 G4Material* vacuum = G4Material::GetMaterial("Vacuum");
101
102
103 //
104 // Shape 0 - Quadrupole 1
105 //
106 G4Material* shape0_mat = nist->FindOrBuildMaterial("G4_Al");
107 G4double shape0_rmin = 3.*mm, shape0_rmax = 1.*cm;
108 G4double shape0_hz = 1.*cm;
109 G4double shape0_phimin = 0.*deg, shape0_phimax = 360.*deg;
110 G4ThreeVector pos0 = G4ThreeVector(0*cm, 0*cm, -20.*cm);
111
112 G4Tubs* solidShape0 =
113 new G4Tubs("Shape0", //its name
114 shape0_rmin, shape0_rmax, shape0_hz, shape0_phimin, shape0_phimax); //its size
115
116 G4LogicalVolume* logicShape0 =
117 new G4LogicalVolume(solidShape0, //its solid
118 shape0_mat, //its material
119 "Shape0"); //its name
120
121 new G4PVPlacement(0, //no rotation
122 pos0, //at position
123 logicShape0, //its logical volume
124 "Shape0", //its name
125 logicEnv, //its mother volume
126 false, //no boolean operation
127 0, //copy number
128 checkOverlaps); //overlaps checking
129
130
131 // Shape 1 - Beam pipe 1
132
133 //Position
134 G4ThreeVector pos1 = pos0;
135
136 G4double shape1_rmin = 0.*mm, shape1_rmax = 3.*mm;
137 G4double shape1_hz = 1.*cm;
```

## Section

```
138 G4double shape1_phimin = 0.*deg, shape1_phimax = 360.*deg;
139 G4Tubs* solidShape1 =
140 new G4Tubs("Shape1",
141 shape1_rmin, shape1_rmax, shape1_hz, shape1_phimin, shape1_phimax);
142
143 flogicShape1 =
144 new G4LogicalVolume(solidShape1,          //its solid
145 vacuum,          //its material
146 "Shape1");          //its name
147
148 new G4PVPlacement(0,          //no rotation
149 pos1,          //at position
150 flogicShape1,          //its logical volume
151 "Shape1",          //its name
152 logicEnv,          //its mother volume
153 false,          //no boolean operation
154 0,          //copy number
155 checkOverlaps);          //overlaps checking
156
157
158 //
159 // Beam pipe envelope
160 //
161 G4Material* shape_mat = nist->FindOrBuildMaterial("G4_STAINLESS-STEEL");
162 G4double shape_rmin = 3.*mm, shape_rmax = 3.2*mm;
163 G4double shape_hz = 3.25*cm;
164 G4double shape_phimin = 0.*deg, shape_phimax = 360.*deg;
165
166 //Position
167 G4ThreeVector pos = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz + shape_hz)
168 ;
169
170 G4Tubs* solidShape =
171 new G4Tubs("Shape",
172 shape_rmin, shape_rmax, shape_hz, shape_phimin, shape_phimax);
173
174 G4LogicalVolume* logicShape =
175 new G4LogicalVolume(solidShape,          //its solid
176 shape_mat,          //its material
177 "Shape");          //its name
178
179 new G4PVPlacement(0,          //no rotation
180 pos,          //at position
181 logicShape,          //its logical volume
182 "Shape",          //its name
183 logicEnv,          //its mother volume
184 false,          //no boolean operation
185 0,          //copy number
186 checkOverlaps          //overlaps checking
187 );
188 //
```

```
189 // Shape 2 - Beam pipe 2
190 //
191 G4double shape2_rmin = 0.*mm, shape2_rmax = 3.*mm;
192 G4double shape2_hz = 3.25*cm;
193 G4double shape2_phimin = 0.*deg, shape2_phimax = 360.*deg;
194
195 //Position
196 G4ThreeVector pos2 = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz +
    shape2_hz);
197
198 G4Tubs* solidShape2 =
199 new G4Tubs("Shape2",
200 shape2_rmin, shape2_rmax, shape2_hz, shape2_phimin, shape2_phimax);
201
202 G4LogicalVolume* logicShape2 =
203 new G4LogicalVolume(solidShape2, //its solid
204 vacuum, //its material
205 "Shape2"); //its name
206
207 new G4PVPlacement(0, //no rotation
208 pos2, //at position
209 logicShape2, //its logical volume
210 "Shape2", //its name
211 logicEnv, //its mother volume
212 false, //no boolean operation
213 0, //copy number
214 checkOverlaps //overlaps checking
215 );
216
217 //Add air layer
218 G4double layer = 0.*mm;
219 //
220 // Shape 3 - Quadrupole 2
221 //
222 G4double shape3_rmin = 3.*mm, shape3_rmax = 1.*cm;
223 G4double shape3_hz = 1.*cm;
224 G4double shape3_phimin = 0.*deg, shape3_phimax = 360.*deg;
225 G4ThreeVector pos3 = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz + 2*
    shape2_hz + layer + shape3_hz);
226 G4Tubs* solidShape3 =
227 new G4Tubs("Shape3",
228 shape3_rmin, shape3_rmax, shape3_hz, shape3_phimin, shape3_phimax);
229
230 G4LogicalVolume* logicShape3 =
231 new G4LogicalVolume(solidShape3, //its solid
232 shape0_mat, //its material
233 "Shape3"); //its name
234
235 new G4PVPlacement(0, //no rotation
236 pos3, //at position
237 logicShape3, //its logical volume
238 "Shape3", //its name
```



## Section

```
239 logicEnv ,           //its mother volume
240 false ,              //no boolean operation
241 0,                   //copy number
242 checkOverlaps);     //overlaps checking
243 //
244 // Shape 4 - Beam pipe 3
245 //
246 G4double shape4_rmin = 0.*mm, shape4_rmax = 3.*mm;
247 G4double shape4_hz = 1.*cm;
248 G4double shape4_phimin = 0.*deg, shape4_phimax = 360.*deg;
249 G4ThreeVector pos4 = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz + 2*
    shape2_hz + layer + shape3_hz);
250 G4Tubs* solidShape4 =
251 new G4Tubs("Shape4",
252 shape4_rmin, shape4_rmax, shape4_hz, shape4_phimin, shape4_phimax);
253
254 flogicShape4 =
255 new G4LogicalVolume(solidShape4,           //its solid
256 vacuum,                                   //its material
257 "Shape4");                                //its name
258
259 new G4PVPlacement(0,                       //no rotation
260 pos4,                                     //at position
261 flogicShape4,                             //its logical volume
262 "Shape4",                                 //its name
263 logicEnv,                                 //its mother volume
264 false,                                    //no boolean operation
265 0,                                        //copy number
266 checkOverlaps);                          //overlaps checking
267
268 //
269 // Shape 5 - Module 0
270 //
271
272 G4Material* shape5_mat = nist->FindOrBuildMaterial("G4_Cu");
273 G4double shape5_rmin = 1.5*mm, shape5_rmax = 3.*cm;
274 G4double shape5_hz = 3.*cm;
275 G4double shape5_phimin = 0.*deg, shape5_phimax = 360.*deg;
276 G4ThreeVector pos5 = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz + 2*
    shape2_hz + layer + 2*shape3_hz + layer + shape5_hz );
277 G4Tubs* solidShape5 =
278 new G4Tubs("Shape5",                       //its name
279 shape5_rmin, shape5_rmax, shape5_hz, shape5_phimin, shape5_phimax); //its size
280
281 G4LogicalVolume* logicShape5 =
282 new G4LogicalVolume(solidShape5,           //its solid
283 shape5_mat,                               //its material
284 "Shape5");                                //its name
285
286 new G4PVPlacement(0,                       //no rotation
287 pos5,                                     //at position
288 logicShape5,                             //its logical volume
```

```
289 "Shape5",           //its name
290 logicEnv ,         //its mother volume
291 false ,           //no boolean operation
292 0,                //copy number
293 checkOverlaps);  //overlaps checking
294 //
295 // Shape 6 - Beam pipe 4
296 //
297 G4double shape6_rmin = 0.*mm, shape6_rmax = 1.5*mm;
298 G4double shape6_hz = 3.*cm;
299 G4double shape6_phimin = 0.*deg, shape6_phimax = 360.*deg;
300 G4ThreeVector pos6 = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz + 2*
    shape2_hz + layer + 2*shape3_hz + layer + shape5_hz);
301 G4Tubs* solidShape6 =
302 new G4Tubs("Shape6",           //its name
303 shape6_rmin, shape6_rmax, shape6_hz, shape6_phimin, shape6_phimax); //its size
304
305 flogicShape6 =
306 new G4LogicalVolume(solidShape6, //its solid
307 vacuum, //its material
308 "Shape6"); //its name
309
310 new G4PVPlacement(0, //no rotation
311 pos6, //at position
312 flogicShape6, //its logical volume
313 "Shape6", //its name
314 logicEnv, //its mother volume
315 false, //no boolean operation
316 0, //copy number
317 checkOverlaps); //overlaps checking
318
319
320 //
321 // Beam pipe between Module 0 and Quadrupole 1
322 //
323 //
324 // Beam pipe envelope
325 //
326 G4Material* beampipe1_mat = nist->FindOrBuildMaterial("G4_STAINLESS-STEEL");
327 G4double beampipe1_rmin = 3.*mm, beampipe1_rmax = 3.2*mm;
328 G4double beampipe1_hz = 0.5*cm;
329 G4double beampipe1_phimin = 0.*deg, beampipe1_phimax = 360.*deg;
330
331 //Position
332 G4ThreeVector beampipe1_pos = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz +
    2*shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + beampipe1_hz);
333
334 G4Tubs* beampipe1_solid =
335 new G4Tubs("beampipe1solid",
336 beampipe1_rmin, beampipe1_rmax, beampipe1_hz, beampipe1_phimin,
    beampipe1_phimax);
337
```

```
338 G4LogicalVolume* beampipe1_logic =
339 new G4LogicalVolume(beampipe1_solid, //its solid
340 beampipe1_mat, //its material
341 "beampipe1logic"); //its name
342
343 new G4PVPlacement(0, //no rotation
344 beampipe1_pos, //at position
345 beampipe1_logic, //its logical volume
346 "beampipe1physical", //its name
347 logicEnv, //its mother volume
348 false, //no boolean operation
349 0, //copy number
350 checkOverlaps //overlaps checking
351 );
352
353 //
354 // Vacuum inside beampipe
355 //
356 G4double vacuum1_rmin = 0.*mm, vacuum1_rmax = 3.*mm;
357 G4double vacuum1_hz = 0.5*cm;
358 G4double vacuum1_phimin = 0.*deg, vacuum1_phimax = 360.*deg;
359
360 //Position
361 G4ThreeVector vacuum1_pos = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz +
    2*shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + vacuum1_hz);
362
363 G4Tubs* vacuum1_solid =
364 new G4Tubs("vacuum1solid",
365 vacuum1_rmin, vacuum1_rmax, vacuum1_hz, vacuum1_phimin, vacuum1_phimax);
366
367 G4LogicalVolume* vacuum1_logic =
368 new G4LogicalVolume(vacuum1_solid, //its solid
369 vacuum, //its material
370 "vacuum1logic"); //its name
371
372 new G4PVPlacement(0, //no rotation
373 vacuum1_pos, //at position
374 vacuum1_logic, //its logical volume
375 "vacuum1physical", //its name
376 logicEnv, //its mother volume
377 false, //no boolean operation
378 0, //copy number
379 checkOverlaps //overlaps checking
380 );
381
382 //
383 // Shape 7 - Quadrupole 3
384 //
385 G4double shape7_rmin = 3.*mm, shape7_rmax = 1.*cm;
386 G4double shape7_hz = 1.*cm;
387 G4double shape7_phimin = 0.*deg, shape7_phimax = 360.*deg;
388 G4ThreeVector pos7 = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz + 2*
```

```
    shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + 2*vacuum1_hz +
    shape7_hz);
389
390 G4Tubs* solidShape7 =
391     new G4Tubs("Shape7",           //its name
392     shape7_rmin, shape7_rmax, shape7_hz, shape7_phimin, shape7_phimax); //its size
393
394 G4LogicalVolume* logicShape7 =
395     new G4LogicalVolume(solidShape7,           //its solid
396     shape0_mat,           //its material
397     "Shape7");           //its name
398
399     new G4PVPlacement(0,           //no rotation
400     pos7,           //at position
401     logicShape7,           //its logical volume
402     "Shape7",           //its name
403     logicEnv,           //its mother volume
404     false,           //no boolean operation
405     0,           //copy number
406     checkOverlaps);           //overlaps checking
407
408     //
409     // Shape 8 - Beam pipe 5
410     //
411     G4double shape8_rmin = 0.*mm, shape8_rmax = 3.*mm;
412     G4double shape8_hz = 1.*cm;
413     G4double shape8_phimin = 0.*deg, shape8_phimax = 360.*deg;
414     G4ThreeVector pos8 = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz + 2*
        shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + 2*vacuum1_hz +
        shape7_hz);
415
416     G4Tubs* solidShape8 =
417     new G4Tubs("Shape8",           //its name
418     shape8_rmin, shape8_rmax, shape8_hz, shape8_phimin, shape8_phimax); //its size
419
420     flogicShape8 =
421     new G4LogicalVolume(solidShape8,           //its solid
422     vacuum,           //its material
423     "Shape8");           //its name
424
425     new G4PVPlacement(0,           //no rotation
426     pos8,           //at position
427     flogicShape8,           //its logical volume
428     "Shape8",           //its name
429     logicEnv,           //its mother volume
430     false,           //no boolean operation
431     0,           //copy number
432     checkOverlaps);           //overlaps checking
433
434     //
435     // Beam pipe between Quadrupole 3 and Module 1
436     //
```

## Section

```
437 //
438 // Beam pipe envelope
439 //
440 G4Material* beampipe2_mat = nist->FindOrBuildMaterial("G4_STAINLESS-STEEL");
441 G4double beampipe2_rmin = 3.*mm, beampipe2_rmax = 3.2*mm;
442 G4double beampipe2_hz = 0.5*cm;
443 G4double beampipe2_phimin = 0.*deg, beampipe2_phimax = 360.*deg;
444
445 //Position
446 G4ThreeVector beampipe2_pos = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz +
      2*shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + 2*vacuum1_hz +
      2*shape7_hz + beampipe2_hz);
447
448 G4Tubs* beampipe2_solid =
449 new G4Tubs("beampipe2solid",
450 beampipe2_rmin, beampipe2_rmax, beampipe2_hz, beampipe2_phimin,
      beampipe2_phimax);
451
452 G4LogicalVolume* beampipe2_logic =
453 new G4LogicalVolume(beampipe2_solid, //its solid
454 beampipe2_mat, //its material
455 "beampipe2logic"); //its name
456
457 new G4PVPlacement(0, //no rotation
458 beampipe2_pos, //at position
459 beampipe2_logic, //its logical volume
460 "beampipe2physical", //its name
461 logicEnv, //its mother volume
462 false, //no boolean operation
463 0, //copy number
464 checkOverlaps //overlaps checking
465 );
466
467 //
468 // Vacuum inside beampipe
469 //
470 G4double vacuum2_rmin = 0.*mm, vacuum2_rmax = 3.*mm;
471 G4double vacuum2_hz = 0.5*cm;
472 G4double vacuum2_phimin = 0.*deg, vacuum2_phimax = 360.*deg;
473
474 //Position
475 G4ThreeVector vacuum2_pos = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz +
      2*shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + 2*vacuum1_hz + 2*
      shape7_hz + vacuum2_hz);
476
477 G4Tubs* vacuum2_solid =
478 new G4Tubs("vacuum2solid",
479 vacuum2_rmin, vacuum2_rmax, vacuum2_hz, vacuum2_phimin, vacuum2_phimax);
480
481 G4LogicalVolume* vacuum2_logic =
482 new G4LogicalVolume(vacuum2_solid, //its solid
483 vacuum, //its material
```

```
484 "vacuum2logic"); //its name
485
486 new G4PVPlacement(0, //no rotation
487 vacuum2_pos, //at position
488 vacuum2_logic, //its logical volume
489 "vacuum2physical", //its name
490 logicEnv, //its mother volume
491 false, //no boolean operation
492 0, //copy number
493 checkOverlaps //overlaps checking
494 );
495
496 // //
497 // -- Modulo 1 --- Quadrupole 3 //
498 // //
499
500 //
501 // Shape 9 - Module 1
502 //
503 G4Material* shape9_mat = nist->FindOrBuildMaterial("G4_Cu");
504 G4double shape9_rmin = 1.5*mm, shape9_rmax = 3.*cm;
505 G4double shape9_hz = 3.*cm;
506 G4double shape9_phimin = 0.*deg, shape9_phimax = 360.*deg;
507 G4ThreeVector pos9 = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz + 2*
    shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + 2*vacuum1_hz + 2*
    shape7_hz + 2*vacuum2_hz + shape9_hz);
508 G4Tubs* solidShape9 =
509 new G4Tubs("Shape9", //its name
510 shape9_rmin, shape9_rmax, shape9_hz, shape9_phimin, shape9_phimax); //its size
511
512 G4LogicalVolume* logicShape9 =
513 new G4LogicalVolume(solidShape9, //its solid
514 shape9_mat, //its material
515 "Shape9"); //its name
516
517 new G4PVPlacement(0, //no rotation
518 pos9, //at position
519 logicShape9, //its logical volume
520 "Shape9", //its name
521 logicEnv, //its mother volume
522 false, //no boolean operation
523 0, //copy number
524 checkOverlaps); //overlaps checking
525 //
526 // Shape 10 - Beam pipe 6
527 //
528 G4double shape10_rmin = 0.*mm, shape10_rmax = 1.5*mm;
529 G4double shape10_hz = 3.*cm;
530 G4double shape10_phimin = 0.*deg, shape10_phimax = 360.*deg;
531 G4ThreeVector pos10 = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz + 2*
    shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + 2*vacuum1_hz + 2*
    shape7_hz + 2*vacuum2_hz + shape9_hz);
```

```
532 G4Tubs* solidShape10 =
533 new G4Tubs("Shape10", //its name
534 shape10_rmin, shape10_rmax, shape10_hz, shape10_phimin, shape10_phimax); //its
    size
535
536 flogicShape10 =
537 new G4LogicalVolume(solidShape10, //its solid
538 vacuum, //its material
539 "Shape10"); //its name
540
541 new G4PVPlacement(0, //no rotation
542 pos10, //at position
543 flogicShape10, //its logical volume
544 "Shape10", //its name
545 logicEnv, //its mother volume
546 false, //no boolean operation
547 0, //copy number
548 checkOverlaps); //overlaps checking
549
550
551 //
552 // Beam pipe between Module 1 and Quadrupole 4
553 //
554 //
555 // Beam pipe envelope
556 //
557 G4Material* beampipe3_mat = nist->FindOrBuildMaterial("G4_STAINLESS-STEEL");
558 G4double beampipe3_rmin = 3.*mm, beampipe3_rmax = 3.2*mm;
559 G4double beampipe3_hz = 0.5*cm;
560 G4double beampipe3_phimin = 0.*deg, beampipe3_phimax = 360.*deg;
561
562 //Position
563 G4ThreeVector beampipe3_pos = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz +
    2*shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + 2*vacuum1_hz +
    2*shape7_hz + 2*beampipe2_hz + 2*shape9_hz + beampipe3_hz);
564
565 G4Tubs* beampipe3_solid =
566 new G4Tubs("beampipe3solid",
567 beampipe3_rmin, beampipe3_rmax, beampipe3_hz, beampipe3_phimin,
    beampipe3_phimax);
568
569 G4LogicalVolume* beampipe3_logic =
570 new G4LogicalVolume(beampipe3_solid, //its solid
571 beampipe3_mat, //its material
572 "beampipe3logic"); //its name
573
574 new G4PVPlacement(0, //no rotation
575 beampipe3_pos, //at position
576 beampipe3_logic, //its logical volume
577 "beampipe3physical", //its name
578 logicEnv, //its mother volume
579 false, //no boolean operation
```

```
580 0, //copy number
581 checkOverlaps //overlaps checking
582 );
583
584 //
585 // Vacuum inside beampipe
586 //
587 G4double vacuum3_rmin = 0.*mm, vacuum3_rmax = 3.*mm;
588 G4double vacuum3_hz = 0.5*cm;
589 G4double vacuum3_phimin = 0.*deg, vacuum3_phimax = 360.*deg;
590
591 //Position
592 G4ThreeVector vacuum3_pos = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz +
    2*shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + 2*vacuum1_hz + 2*
    shape7_hz + 2*vacuum2_hz + 2*shape9_hz + vacuum3_hz);
593
594 G4Tubs* vacuum3_solid =
595 new G4Tubs("vacuum3solid",
596 vacuum3_rmin, vacuum3_rmax, vacuum3_hz, vacuum3_phimin, vacuum3_phimax);
597
598 G4LogicalVolume* vacuum3_logic =
599 new G4LogicalVolume(vacuum3_solid, //its solid
600 vacuum, //its material
601 "vacuum3logic"); //its name
602
603 new G4PVPlacement(0, //no rotation
604 vacuum3_pos, //at position
605 vacuum3_logic, //its logical volume
606 "vacuum3physical", //its name
607 logicEnv, //its mother volume
608 false, //no boolean operation
609 0, //copy number
610 checkOverlaps //overlaps checking
611 );
612
613 //
614 // Shape 11 - Quadrupole 4
615 //
616 G4double shape11_rmin = 3.*mm, shape11_rmax = 1.*cm;
617 G4double shape11_hz = 1.*cm;
618 G4double shape11_phimin = 0.*deg, shape11_phimax = 360.*deg;
619 G4ThreeVector pos11 = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz + 2*
    shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + 2*vacuum1_hz + 2*
    shape7_hz + 2*vacuum2_hz + 2*shape9_hz + 2*vacuum3_hz + shape11_hz);
620
621 G4Tubs* solidShape11 =
622 new G4Tubs("Shape11", //its name
623 shape11_rmin, shape11_rmax, shape11_hz, shape11_phimin, shape11_phimax); //its
    size
624
625 G4LogicalVolume* logicShape11 =
626 new G4LogicalVolume(solidShape11, //its solid
```



## Section

---

```

627 shape0_mat,          //its material
628 "Shape11");         //its name
629
630 new G4PVPlacement(0,          //no rotation
631 pos11,                //at position
632 logicShape11,         //its logical volume
633 "Shape11",           //its name
634 logicEnv,            //its mother volume
635 false,               //no boolean operation
636 0,                   //copy number
637 checkOverlaps);     //overlaps checking
638
639 //
640 // Shape 12 - Beam pipe 7
641 //
642 G4double shape12_rmin = 0.*mm, shape12_rmax = 3.*mm;
643 G4double shape12_hz = 1.*cm;
644 G4double shape12_phimin = 0.*deg, shape12_phimax = 360.*deg;
645 G4ThreeVector pos12 = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz + 2*
        shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + 2*vacuum1_hz + 2*
        shape7_hz + 2*vacuum2_hz + 2*shape9_hz + 2*vacuum3_hz + shape11_hz);
646
647 G4Tubs* solidShape12 =
648 new G4Tubs("Shape12",          //its name
649 shape12_rmin, shape12_rmax, shape12_hz, shape12_phimin, shape12_phimax); //its
        size
650
651 flogicShape12 =
652 new G4LogicalVolume(solidShape12, //its solid
653 vacuum,           //its material
654 "Shape12");       //its name
655
656 new G4PVPlacement(0,          //no rotation
657 pos12,                //at position
658 flogicShape12,         //its logical volume
659 "Shape12",           //its name
660 logicEnv,            //its mother volume
661 false,               //no boolean operation
662 0,                   //copy number
663 checkOverlaps);     //overlaps checking
664
665 //
666 // Beam pipe between Quadrupole 4 and Module 2
667 //
668 //
669 // Beam pipe envelope
670 //
671 G4Material* beampipe4_mat = nist->FindOrBuildMaterial("G4_STAINLESS-STEEL");
672 G4double beampipe4_rmin = 3.*mm, beampipe4_rmax = 3.2*mm;
673 G4double beampipe4_hz = 0.5*cm;
674 G4double beampipe4_phimin = 0.*deg, beampipe4_phimax = 360.*deg;
675

```

```
676 //Position
677 G4ThreeVector beampipe4_pos = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz +
    2*shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + 2*vacuum1_hz +
    2*shape7_hz + 2*beampipe2_hz + 2*shape9_hz + 2*beampipe3_hz + 2*shape11_hz
    + beampipe4_hz);
678
679 G4Tubs* beampipe4_solid =
680 new G4Tubs("beampipe4solid",
681 beampipe4_rmin, beampipe4_rmax, beampipe4_hz, beampipe4_phimin,
    beampipe4_phimax);
682
683 G4LogicalVolume* beampipe4_logic =
684 new G4LogicalVolume(beampipe4_solid, //its solid
685 beampipe4_mat, //its material
686 "beampipe4logic"); //its name
687
688 new G4PVPlacement(0, //no rotation
689 beampipe4_pos, //at position
690 beampipe4_logic, //its logical volume
691 "beampipe4physical", //its name
692 logicEnv, //its mother volume
693 false, //no boolean operation
694 0, //copy number
695 checkOverlaps //overlaps checking
696 );
697
698 //
699 // Vacuum inside beampipe
700 //
701 G4double vacuum4_rmin = 0.*mm, vacuum4_rmax = 3.*mm;
702 G4double vacuum4_hz = 0.5*cm;
703 G4double vacuum4_phimin = 0.*deg, vacuum4_phimax = 360.*deg;
704
705 //Position
706 G4ThreeVector vacuum4_pos = G4ThreeVector(0*cm, 0*cm, -20.*cm + shape1_hz +
    2*shape2_hz + layer + 2*shape3_hz + layer + 2*shape5_hz + 2*vacuum1_hz + 2*
    shape7_hz + 2*vacuum2_hz + 2*shape9_hz + 2*vacuum3_hz + 2*shape11_hz +
    beampipe4_hz);
707
708 G4Tubs* vacuum4_solid =
709 new G4Tubs("vacuum4solid",
710 vacuum4_rmin, vacuum4_rmax, vacuum4_hz, vacuum4_phimin, vacuum4_phimax);
711
712 G4LogicalVolume* vacuum4_logic =
713 new G4LogicalVolume(vacuum4_solid, //its solid
714 vacuum, //its material
715 "vacuum4logic"); //its name
716
717 new G4PVPlacement(0, //no rotation
718 vacuum4_pos, //at position
719 vacuum4_logic, //its logical volume
720 "vacuum4physical", //its name
```

## Section

```
721 logicEnv ,           //its mother volume
722 false ,             //no boolean operation
723 0,                  //copy number
724 checkOverlaps       //overlaps checking
725 );
726
727
728 //
729 // Construct Assembled Volume
730 //
731 G4AssemblyVolume* assembly = new G4AssemblyVolume();
732
733 assembly->AddPlacedVolume( logicShape5 ,      pos5 ,      0);
734 assembly->AddPlacedVolume( flogicShape6 ,     pos6 ,      0);
735 assembly->AddPlacedVolume( beampipe1_logic ,  beampipe1_pos , 0);
736 assembly->AddPlacedVolume( vacuum1_logic ,   vacuum1_pos ,  0);
737 assembly->AddPlacedVolume( logicShape7 ,     pos7 ,      0);
738 assembly->AddPlacedVolume( flogicShape8 ,     pos8 ,      0);
739 assembly->AddPlacedVolume( beampipe2_logic ,  beampipe2_pos , 0);
740 assembly->AddPlacedVolume( vacuum2_logic ,   vacuum2_pos ,  0);
741 assembly->AddPlacedVolume( logicShape9 ,     pos9 ,      0);
742 assembly->AddPlacedVolume( flogicShape10 ,   pos10 ,     0);
743 assembly->AddPlacedVolume( beampipe3_logic ,  beampipe3_pos , 0);
744 assembly->AddPlacedVolume( vacuum3_logic ,   vacuum3_pos ,  0);
745 assembly->AddPlacedVolume( logicShape11 ,    pos11 ,     0);
746 assembly->AddPlacedVolume( flogicShape12 ,   pos12 ,     0);
747 assembly->AddPlacedVolume( beampipe4_logic ,  beampipe4_pos , 0);
748 assembly->AddPlacedVolume( vacuum4_logic ,   vacuum4_pos ,  0);
749
750 G4double z_pos = 2*shape5_hz + 2*vacuum1_hz + 2*shape7_hz + 2*vacuum2_hz + 2*
    shape9_hz + 2*vacuum3_hz + 2*shape11_hz + 2*beampipe4_hz;
751
752 for (int i = 0; i < 4; i++)
753 {
754     G4ThreeVector position = G4ThreeVector(0*cm, 0*cm, z_pos*(i+1));
755     assembly->MakeImprint(logicEnv , position , 0);
756 }
757
758 SetLogicalVolume(logicEnv);
759 }
760
761 // .....oooOOOOooo ..... oooOOOOooo ..... oooOOOOooo ..... oooOOOOooo
    .....
```

## Appendix B

# LXeDetectorConstruction

GEANT4, C++ LXeDetectorConstruction.cc class:

```
1 //
2 // *****
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4 // *
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23 // * acceptance of all terms of the Geant4 Software license. *
24 // *****
25 //
26 //
27 /// \file optical/LXe/src/LXeDetectorConstruction.cc
28 /// \brief Implementation of the LXeDetectorConstruction class
29 //
30 //
31 #include "LXeDetectorConstruction.hh"
32
33 #include "LXeDetectorMessenger.hh"
34 #include "LXeMainVolume.hh"
35 #include "LXePMTSD.hh"
```

## Section

---

```
36 #include "LXeScintSD.hh"
37 #include "LXeWLSSlab.hh"
38 #include "LXeAccelerator.hh"
39 #include "LXeModule1.hh"
40 #include "LXeModule2.hh"
41 #include "LXeModule3.hh"
42 #include "LXeModuleCCL.hh"
43
44 #include "G4Types.hh"
45 #include "ElectricFieldSetup.hh"
46 #include "F03FieldSetup.hh"
47
48 #include "LXeSteppingAction.hh"
49
50 #include "globals.hh"
51 #include "G4Box.hh"
52 #include "G4GeometryManager.hh"
53 #include "G4LogicalBorderSurface.hh"
54 #include "G4LogicalSkinSurface.hh"
55 #include "G4LogicalVolume.hh"
56 #include "G4LogicalVolumeStore.hh"
57 #include "G4Material.hh"
58 #include "G4MaterialTable.hh"
59 #include "G4NistManager.hh"
60 #include "G4OpticalSurface.hh"
61 #include "G4PhysicalConstants.hh"
62 #include "G4PhysicalVolumeStore.hh"
63 #include "G4PVPlacement.hh"
64 #include "G4RunManager.hh"
65 #include "G4SDManager.hh"
66 #include "G4SolidStore.hh"
67 #include "G4Sphere.hh"
68 #include "G4SystemOfUnits.hh"
69 #include "G4ThreeVector.hh"
70 #include "G4Tubs.hh"
71 #include "G4UImanager.hh"
72 #include "G4VisAttributes.hh"
73 #include "G4AutoDelete.hh"
74
75 using namespace CLHEP;
76
77 G4bool LXeDetectorConstruction::fSphereOn = true;
78
79 // .....oooOOOOooo ..... oooOOOOooo ..... oooOOOOooo ..... oooOOOOooo
80 .....
81 LXeDetectorConstruction::LXeDetectorConstruction()
82 : fLXe_mt(nullptr)
83 , fMPTPStyrene(nullptr)
84 , fScoringVolume(nullptr)
85 , fMPTGlassSci(nullptr)
86 {
```

```
87 fExperimentalHall_box = nullptr;
88 fExperimentalHall_log = nullptr;
89 fExperimentalHall_phys = nullptr;
90
91 fLXe = fAl = fAir = fVacuum = fGlass = fGlassSci = nullptr;
92 fPstyrene = fPMMA = fPethylene1 = fPethylene2 = nullptr;
93 fConcrete = nullptr;
94
95 fN = fO = fC = fH = fLi6 = fCe = nullptr;
96 fAll = nullptr;
97 fNa = fMg = fSi = fK = fCa = fFe = fP = fS = fTi = fMn = fZn = fZr = fBa =
    fPb = fSr = nullptr;
98
99 fSaveThreshold = 0;
100 SetDefaults();
101
102 DefineMaterials();
103 fDetectorMessenger = new LXeDetectorMessenger(this);
104
105 felFieldSetup1_0 = 0;
106 felFieldSetup2_0 = 0;
107 femFieldSetup1_0 = 0;
108 femFieldSetup2_0 = 0;
109 femFieldSetup3_0 = 0;
110 femFieldSetup4_0 = 0;
111
112 felFieldSetup1_1 = 0;
113 felFieldSetup2_1 = 0;
114 femFieldSetup1_1 = 0;
115 femFieldSetup2_1 = 0;
116
117 felFieldSetup1_2 = 0;
118 felFieldSetup2_2 = 0;
119 femFieldSetup1_2 = 0;
120 femFieldSetup2_2 = 0;
121
122 felFieldSetup1_3 = 0;
123 felFieldSetup2_3 = 0;
124 femFieldSetup1_3 = 0;
125 femFieldSetup2_3 = 0;
126
127 felFieldSetup1_4 = 0;
128 felFieldSetup2_4 = 0;
129 femFieldSetup1_4 = 0;
130 femFieldSetup2_4 = 0;
131
132 felFieldSetup1_5 = 0;
133 felFieldSetup2_5 = 0;
134 femFieldSetup1_5 = 0;
135 femFieldSetup2_5 = 0;
136
137 felFieldSetup1_6 = 0;
```

## Section

---

```

138 felFieldSetup2_6 = 0;
139 femFieldSetup1_6 = 0;
140 femFieldSetup2_6 = 0;
141
142 felFieldSetup1_7 = 0;
143 felFieldSetup2_7 = 0;
144 femFieldSetup1_7 = 0;
145 femFieldSetup2_7 = 0;
146 }
147
148 // .....oooOO0OOooo ..... oooOO0OOooo ..... oooOO0OOooo ..... oooOO0OOooo
149 .....
150 LXeDetectorConstruction::~LXeDetectorConstruction()
151 {
152     if (fMainVolume)
153     {
154         delete fMainVolume;
155     }
156     delete fLXe_mt;
157     delete fDetectorMessenger;
158     delete fMPTPStyrene;
159     delete fScoringVolume;
160     delete fMPTGlassSci;
161 }
162
163 // .....oooOO0OOooo ..... oooOO0OOooo ..... oooOO0OOooo ..... oooOO0OOooo
164 .....
165 void LXeDetectorConstruction::DefineMaterials()
166 {
167     G4double a; // atomic mass
168     G4double z; // atomic number
169     G4double density;
170
171     G4int polyPMMA = 1;
172     G4int nC_PMMA = 3 + 2 * polyPMMA;
173     G4int nH_PMMA = 6 + 2 * polyPMMA;
174
175     G4int polyeth = 1;
176     G4int nC_eth = 2 * polyeth;
177     G4int nH_eth = 4 * polyeth;
178
179     /*** Elements
180     fH = new G4Element("H", "H", z = 1., a = 1.01 * g / mole);
181     fC = new G4Element("C", "C", z = 6., a = 12.01 * g / mole);
182     fN = new G4Element("N", "N", z = 7., a = 14.01 * g / mole);
183     fO = new G4Element("O", "O", z = 8., a = 16.00 * g / mole);
184     fLi6 = new G4Element("Li6", "Li6", z = 3., a = 6.00 * g / mole);
185     fCe = new G4Element("Ce", "Ce", z = 58., a = 140.12 * g / mole);
186
187     fAl = new G4Element("Al", "Al", z = 13., a = 26.98 * g / mole);

```

```
188
189 fNa = new G4Element("Na", "Na", z = 11., a = 22.99 * g / mole);
190 fMg = new G4Element("Mg", "Mg", z = 12., a = 24.30 * g / mole);
191 fSi = new G4Element("Si", "Si", z = 14., a = 28.08 * g / mole);
192 fK = new G4Element("K", "K", z = 19., a = 39.10 * g / mole);
193 fCa = new G4Element("Ca", "Ca", z = 20., a = 40.078 * g / mole);
194 fFe = new G4Element("Fe", "Fe", z = 26., a = 55.84 * g / mole);
195 fP = new G4Element("P", "P", z = 15., a = 30.97 * g / mole);
196 fS = new G4Element("S", "S", z = 16., a = 32.065 * g / mole);
197 fTi = new G4Element("Ti", "Ti", z = 22., a = 47.867 * g / mole);
198 fMn = new G4Element("Mn", "Mn", z = 25., a = 54.94 * g / mole);
199 fZn = new G4Element("Zn", "Zn", z = 30., a = 65.409 * g / mole);
200 fZr = new G4Element("Zr", "Zr", z = 40., a = 91.224 * g / mole);
201 fBa = new G4Element("Ba", "Ba", z = 56., a = 137.327 * g / mole);
202 fPb = new G4Element("Pb", "Pb", z = 82., a = 207.2 * g / mole);
203 fSr = new G4Element("Sr", "Sr", z = 38., a = 87.62 * g / mole);
204
205 //*** Materials
206 // Liquid Xenon
207 fLXe = new G4Material("LXe", z = 54., a = 131.29 * g / mole,
208 density = 3.020 * g / cm3);
209 // Aluminum
210 fAl = new G4Material("Al", z = 13., a = 26.98 * g / mole,
211 density = 2.7 * g / cm3);
212 // Vacuum
213 fVacuum = new G4Material("Vacuum", density = universe_mean_density , 3,
    kStateGas ,
214 0.1 * kelvin , 1.e-7 * pascal);
215 fVacuum->AddElement(fH, 1. * perCent);
216 fVacuum->AddElement(fN, 69. * perCent);
217 fVacuum->AddElement(fO, 30. * perCent);
218 // Air
219 fAir = new G4Material("Air", density = 1.29 * mg / cm3, 3);
220 fAir->AddElement(fH, 1. * perCent);
221 fAir->AddElement(fN, 69. * perCent);
222 fAir->AddElement(fO, 30. * perCent);
223 // Glass
224 fGlass = new G4Material("Glass", density = 1.032 * g / cm3, 2);
225 fGlass->AddElement(fC, 91.533 * perCent);
226 fGlass->AddElement(fH, 8.467 * perCent);
227 // Glass scintillator
228 fGlassSci = new G4Material("GlassSci", density = 2.40 * g / cm3, 4);
229 fGlassSci->AddElement(fC, 84.18 * perCent);
230 fGlassSci->AddElement(fH, 7.82 * perCent);
231 fGlassSci->AddElement(fLi6, 7.5 * perCent);
232 fGlassSci->AddElement(fCe, 0.5 * perCent);
233 // Polystyrene
234 fPstyrene = new G4Material("Polystyrene", density = 1.03 * g / cm3, 2);
235 fPstyrene->AddElement(fC, 8);
236 fPstyrene->AddElement(fH, 8);
237 // Fiber (PMMA)
238 fPMMA = new G4Material("PMMA", density = 1190. * kg / m3, 3);
```



```
239 fPMMA->AddElement(fH, nH_PMMA);
240 fPMMA->AddElement(fC, nC_PMMA);
241 fPMMA->AddElement(fO, 2);
242 // Cladding(polyethylene)
243 fPethylene1 = new G4Material("Pethylene1", density = 1200. * kg / m3, 2);
244 fPethylene1->AddElement(fH, nH_eth);
245 fPethylene1->AddElement(fC, nC_eth);
246 // Double cladding(flourinated polyethylene)
247 fPethylene2 = new G4Material("Pethylene2", density = 1400. * kg / m3, 2);
248 fPethylene2->AddElement(fH, nH_eth);
249 fPethylene2->AddElement(fC, nC_eth);
250 // Concrete composition
251 fConcrete = new G4Material("Concrete", density = 2.42 * g / cm3, 19);
252 fConcrete->AddElement(fO, 49.2875 * perCent);
253 fConcrete->AddElement(fC, 5.62 * perCent);
254 fConcrete->AddElement(fH, 0.6 * perCent);
255 fConcrete->AddElement(fNa, 0.453 * perCent);
256 fConcrete->AddElement(fMg, 0.663 * perCent);
257 fConcrete->AddElement(fAl, 2.063 * perCent);
258 fConcrete->AddElement(fSi, 18.867 * perCent);
259 fConcrete->AddElement(fK, 0.656 * perCent);
260 fConcrete->AddElement(fCa, 20.091 * perCent);
261 fConcrete->AddElement(fFe, 1.118 * perCent);
262 fConcrete->AddElement(fP, 0.048 * perCent);
263 fConcrete->AddElement(fS, 0.012 * perCent);
264 fConcrete->AddElement(fTi, 0.347 * perCent);
265 fConcrete->AddElement(fMn, 0.0387 * perCent);
266 fConcrete->AddElement(fZn, 0.0241 * perCent);
267 fConcrete->AddElement(fZr, 0.0074 * perCent);
268 fConcrete->AddElement(fBa, 0.0179 * perCent);
269 fConcrete->AddElement(fPb, 0.0464 * perCent);
270 fConcrete->AddElement(fSr, 0.04 * perCent);
271
272 //***Material properties tables
273
274 std::vector<G4double> lxe_Energy = { 7.0 * eV, 7.07 * eV, 7.14 * eV };
275
276 std::vector<G4double> lxe_SCINT = { 0.1, 1.0, 0.1 };
277 std::vector<G4double> lxe_RIND = { 1.59, 1.57, 1.54 };
278 std::vector<G4double> lxe_ABSL = { 35. * cm, 35. * cm, 35. * cm };
279 fLXe_mt = new G4MaterialPropertiesTable();
280 fLXe_mt->AddProperty("SCINTILLATIONCOMPONENT1", lxe_Energy, lxe_SCINT);
281 fLXe_mt->AddProperty("SCINTILLATIONCOMPONENT2", lxe_Energy, lxe_SCINT);
282 fLXe_mt->AddProperty("RINDEX", lxe_Energy, lxe_RIND);
283 fLXe_mt->AddProperty("ABSLLENGTH", lxe_Energy, lxe_ABSL);
284 fLXe_mt->AddConstProperty("SCINTILLATIONYIELD", 12000. / MeV);
285 fLXe_mt->AddConstProperty("RESOLUTIONSCALE", 1.0);
286 fLXe_mt->AddConstProperty("SCINTILLATIONTIMECONSTANT1", 20. * ns);
287 fLXe_mt->AddConstProperty("SCINTILLATIONTIMECONSTANT2", 45. * ns);
288 fLXe_mt->AddConstProperty("SCINTILLATIONYIELD1", 1.0);
289 fLXe_mt->AddConstProperty("SCINTILLATIONYIELD2", 0.0);
290 fLXe->SetMaterialPropertiesTable(fLXe_mt);
```

```
291
292 // Set the Birks Constant for the LXe scintillator
293 fLXe->GetIonisation()->SetBirksConstant(0.126 * mm / MeV);
294
295 std::vector<G4double> glass_RIND      = { 1.49, 1.49, 1.49 };
296 std::vector<G4double> glass_AbsLength = { 420. * cm, 420. * cm, 420. * cm };
297 G4MaterialPropertiesTable* glass_mt   = new G4MaterialPropertiesTable();
298 glass_mt->AddProperty("ABSLENGTH", lxe_Energy, glass_AbsLength);
299 glass_mt->AddProperty("RINDEX", lxe_Energy, glass_RIND);
300 fGlass->SetMaterialPropertiesTable(glass_mt);
301
302 std::vector<G4double> vacuum_Energy  = { 2.0 * eV, 7.0 * eV, 7.14 * eV };
303 std::vector<G4double> vacuum_RIND    = { 1., 1., 1. };
304 G4MaterialPropertiesTable* vacuum_mt = new G4MaterialPropertiesTable();
305 vacuum_mt->AddProperty("RINDEX", vacuum_Energy, vacuum_RIND);
306 fVacuum->SetMaterialPropertiesTable(vacuum_mt);
307 fAir->SetMaterialPropertiesTable(vacuum_mt); // Give air the same rindex
308
309 std::vector<G4double> wls_Energy = { 2.00 * eV, 2.87 * eV, 2.90 * eV,
310   3.47 * eV };
311
312 std::vector<G4double> rIndexPstyrene = { 1.58, 1.58, 1.58, 1.58 };
313 std::vector<G4double> absorption1    = { 250. * cm, 250. * cm, 250. * cm,
314   250. * cm };
315
316 std::vector<G4double> scintilFast    = { 0.0, 0.0, 1.0, 1.0 };
317 fMPTPStyrene = new G4MaterialPropertiesTable();
318 fMPTPStyrene->AddProperty("RINDEX", wls_Energy, rIndexPstyrene);
319 fMPTPStyrene->AddProperty("ABSLENGTH", wls_Energy, absorption1);
320 fMPTPStyrene->AddProperty("SCINTILLATIONCOMPONENT1", wls_Energy, scintilFast);
321 ;
322 fMPTPStyrene->AddConstProperty("SCINTILLATIONYIELD", 10. / keV);
323 fMPTPStyrene->AddConstProperty("RESOLUTIONSCALE", 1.0);
324 fMPTPStyrene->AddConstProperty("SCINTILLATIONTIMECONSTANT1", 2.4 * ns);
325 fPStyrene->SetMaterialPropertiesTable(fMPTPStyrene);
326
327 // Set the Birks Constant for the Polystyrene scintillator
328 fPStyrene->GetIonisation()->SetBirksConstant(0.126 * mm / MeV);
329
330 //New Material
331 std::vector<G4double> rIndex_GlassSci = { 1.59, 1.59, 1.59, 1.59 };
332 std::vector<G4double> AbsLength_GlassSci = { 420. * cm, 420. * cm, 420. *
333   cm, 420. * cm };
334 std::vector<G4double> scintilFast_GlassSci = { 0.0, 0.0, 1.0, 1.0 };
335 fMPTGlassSci = new G4MaterialPropertiesTable();
336 fMPTGlassSci->AddProperty("RINDEX", wls_Energy, rIndex_GlassSci);
337 fMPTGlassSci->AddProperty("ABSLENGTH", wls_Energy, AbsLength_GlassSci);
338 fMPTGlassSci->AddProperty("SCINTILLATIONCOMPONENT1", wls_Energy,
339   scintilFast_GlassSci);
340 fMPTGlassSci->AddProperty("SCINTILLATIONCOMPONENT2", wls_Energy,
341   scintilFast_GlassSci);
342 fMPTGlassSci->AddConstProperty("SCINTILLATIONYIELD", 10000. / MeV);
343 fMPTGlassSci->AddConstProperty("RESOLUTIONSCALE", 1.0);
```

## Section

---

```

338 fMPTGlassSci->AddConstProperty("SCINTILLATIONTIMECONSTANT1", 18. * ns);
339 fMPTGlassSci->AddConstProperty("SCINTILLATIONTIMECONSTANT2", 45. * ns);
340 fMPTGlassSci->AddConstProperty("SCINTILLATIONYIELD1", 1.);
341 fMPTGlassSci->AddConstProperty("SCINTILLATIONYIELD2", 0.);
342 fGlassSci->SetMaterialPropertiesTable(fMPTGlassSci);
343
344 // Set the Birks Constant for the GlassSci scintillator
345 fGlassSci->GetIonisation()->SetBirksConstant(0.126 * mm / MeV);
346
347
348 std::vector<G4double> RefractiveIndexFiber = { 1.6, 1.6, 1.6, 1.6 };
349 std::vector<G4double> AbsFiber      = { 9.0 * m, 9.0 * m, 0.1 * mm, 0.1 * mm };
350 std::vector<G4double> EmissionFib = { 1.0, 1.0, 0.0, 0.0 };
351 G4MaterialPropertiesTable* fiberProperty = new G4MaterialPropertiesTable();
352 fiberProperty->AddProperty("RINDEX", wls_Energy, RefractiveIndexFiber);
353 fiberProperty->AddProperty("WLSABSLLENGTH", wls_Energy, AbsFiber);
354 fiberProperty->AddProperty("WLSCOMPONENT", wls_Energy, EmissionFib);
355 fiberProperty->AddConstProperty("WLSTIMECONSTANT", 0.5 * ns);
356 fPMMA->SetMaterialPropertiesTable(fiberProperty);
357
358 std::vector<G4double> RefractiveIndexClad1 = { 1.49, 1.49, 1.49, 1.49 };
359 G4MaterialPropertiesTable* clad1Property  = new G4MaterialPropertiesTable();
360 clad1Property->AddProperty("RINDEX", wls_Energy, RefractiveIndexClad1);
361 clad1Property->AddProperty("ABSLLENGTH", wls_Energy, AbsFiber);
362 fPethylene1->SetMaterialPropertiesTable(clad1Property);
363
364 std::vector<G4double> RefractiveIndexClad2 = { 1.42, 1.42, 1.42, 1.42 };
365 G4MaterialPropertiesTable* clad2Property  = new G4MaterialPropertiesTable();
366 clad2Property->AddProperty("RINDEX", wls_Energy, RefractiveIndexClad2);
367 clad2Property->AddProperty("ABSLLENGTH", wls_Energy, AbsFiber);
368 fPethylene2->SetMaterialPropertiesTable(clad2Property);
369 }
370
371 // .....oooOOOOooo ..... oooOOOOooo ..... oooOOOOooo ..... oooOOOOooo
372     .....
373 G4VPhysicalVolume* LXeDetectorConstruction::Construct()
374 {
375     // The experimental hall walls are all 1m away from housing walls
376     G4double expHall_x = fScint_x + fD_mtl + 1.1 * m;
377     G4double expHall_y = fScint_y + fD_mtl + 1.1 * m;
378     G4double expHall_z = ((1.79 + 6. + 4.*1.31 + 0.67 + 3.2) / 4.)*m;
379
380     G4double wall_x = expHall_x + 1.20*m;
381     G4double wall_y = expHall_y + 1.20*m;
382     G4double wall_z = expHall_z + 1.20*m;
383
384     //Create wall over experimental hall
385     fWall_box =
386     new G4Box("wall_box", wall_x, wall_y, wall_z);
387     fWall_log =
388     new G4LogicalVolume(fWall_box, fConcrete, "wall_log", 0, 0, 0);

```

```
389 fWall_phys =
390 new G4PVPlacement(0, G4ThreeVector(), fWall_log, "wall", 0, false, 0);
391
392 //Create experimental hall
393 fExperimentalHall_box =
394 new G4Box("expHall_box", expHall_x, expHall_y, expHall_z);
395 fExperimentalHall_log =
396 new G4LogicalVolume(fExperimentalHall_box, fAir, "expHall_log", 0, 0, 0);
397 fExperimentalHall_phys =
398 new G4PVPlacement(0, G4ThreeVector(), fExperimentalHall_log, "expHall",
    fWall_log, false, 0);
399
400 fExperimentalHall_log->SetVisAttributes(G4Colour(0.8, 0.8, 0.8));
401 fWall_log->SetVisAttributes(G4Colour(0.8, 0.8, 0.8));
402
403 // Place the main volume
404 if (fMainVolumeOn)
405 {
406     fMainVolume = new LXeMainVolume(0, G4ThreeVector(1. *m, 0. *m, -6.60 *m +
        6.0*m), fExperimentalHall_log, false, 0, this);
407 }
408
409 fLXeAccelerator = static_cast<const LXeAccelerator*>(new LXeAccelerator(0,
    G4ThreeVector( 0. , 0. , -6.60*m + 5.0*m), fExperimentalHall_log, false, 0,
    this));
410
411 fLXeModule1 = static_cast<const LXeModule1*>(new LXeModule1(0, G4ThreeVector(
    0. , 0. , 0.395*m - 5.60*m + 5.0*m), fExperimentalHall_log, false, 0, this
    ));
412
413 fLXeModule2 = static_cast<const LXeModule2*>(new LXeModule2(0, G4ThreeVector(
    0. , 0. , 0.395*m - 4.6*m + 5.0*m), fExperimentalHall_log, false, 0, this)
    );
414
415 fLXeModule3 = static_cast<const LXeModule3*>(new LXeModule3(0, G4ThreeVector(
    0. , 0. , 1.395*m - 4.6*m + 5.0*m), fExperimentalHall_log, false, 0, this)
    );
416
417 fLXeModuleCCL1 = static_cast<const LXeModuleCCL*>(new LXeModuleCCL(0,
    G4ThreeVector( 0. , 0. , 2.2225*m - 4.6*m + 5.0*m), fExperimentalHall_log ,
    false, 0, this));
418
419 fLXeModuleCCL2 = static_cast<const LXeModuleCCL*>(new LXeModuleCCL(0,
    G4ThreeVector( 0. , 0. , 2.8775*m - 4.6*m + 5.0*m), fExperimentalHall_log ,
    false, 0, this));
420
421 fLXeModuleCCL3 = static_cast<const LXeModuleCCL*>(new LXeModuleCCL(0,
    G4ThreeVector( 0. , 0. , 3.5325*m - 4.6*m + 5.0*m), fExperimentalHall_log ,
    false, 0, this));
422
423 fLXeModuleCCL4 = static_cast<const LXeModuleCCL*>(new LXeModuleCCL(0,
    G4ThreeVector( 0. , 0. , 4.1875*m - 4.6*m + 5.0*m), fExperimentalHall_log ,
```

```
    false , 0, this));
424
425 // Place the WLS slab
426 if (fWLSslab)
427 {
428     G4VPhysicalVolume* slab = new LXeWLSslab(
429     0, G4ThreeVector(0., 0., -fScint_z / 2. - fSlab_z - 1. * cm),
430     fExperimentalHall_log, false, 0, this);
431
432 // Surface properties for the WLS slab
433 G4OpticalSurface* scintWrap = new G4OpticalSurface("ScintWrap");
434
435 new G4LogicalBorderSurface("ScintWrap", slab, fExperimentalHall_phys,
436 scintWrap);
437
438 scintWrap->SetType(dielectric_metal);
439 scintWrap->SetFinish(polished);
440 scintWrap->SetModel(glisur);
441
442 std::vector<G4double> pp          = { 2.0 * eV, 3.5 * eV };
443 std::vector<G4double> reflectivity = { 1.0, 1.0 };
444 std::vector<G4double> efficiency  = { 0.0, 0.0 };
445
446 G4MaterialPropertiesTable* scintWrapProperty =
447 new G4MaterialPropertiesTable();
448
449 scintWrapProperty->AddProperty("REFLECTIVITY", pp, reflectivity);
450 scintWrapProperty->AddProperty("EFFICIENCY", pp, efficiency);
451 scintWrap->SetMaterialPropertiesTable(scintWrapProperty);
452 }
453
454 fScoringVolume = fExperimentalHall_log;
455
456 return fWall_phys;
457 }
458
459 // .....oooOOOOOooo ..... oooOOOOOooo ..... oooOOOOOooo ..... oooOOOOOooo
460 .....
461 void LXeDetectorConstruction::ConstructSDandField()
462 {
463     if (!fMainVolume)
464         return;
465
466     // PMT SD
467
468     LXePMTSD* pmt = fPmt_SD.Get();
469     if (!pmt)
470     {
471         // Created here so it exists as pmts are being placed
472         G4cout << "Construction /LXeDet/pmtSD" << G4endl;
473         LXePMTSD* pmt_SD = new LXePMTSD("/LXeDet/pmtSD");
474     }
475 }
```

```
474     fPmt_SD.Put(pmt_SD);
475
476     pmt_SD->InitPMTs();
477     pmt_SD->SetPmtPositions(fMainVolume->GetPmtPositions());
478 }
479 else
480 {
481     pmt->InitPMTs();
482     pmt->SetPmtPositions(fMainVolume->GetPmtPositions());
483 }
484 G4SDManager::GetSDMpointer()->AddNewDetector(fPmt_SD.Get());
485 // sensitive detector is not actually on the photocathode.
486 // processHits gets done manually by the stepping action.
487 // It is used to detect when photons hit and get absorbed & detected at the
488 // boundary to the photocathode (which doesn't get done by attaching it to a
489 // logical volume.
490 // It does however need to be attached to something or else it doesn't get
491 // reset at the beginning of events
492
493 SetSensitiveDetector(fMainVolume->GetLogPhotoCath(), fPmt_SD.Get());
494
495 // Scint SD
496
497 if(!fScint_SD.Get())
498 {
499     G4cout << "Construction /LXeDet/scintSD" << G4endl;
500     LXeScintSD* scint_SD = new LXeScintSD("/LXeDet/scintSD");
501     fScint_SD.Put(scint_SD);
502 }
503 G4SDManager::GetSDMpointer()->AddNewDetector(fScint_SD.Get());
504 SetSensitiveDetector(fMainVolume->GetLogScint(), fScint_SD.Get());
505
506 // LXeAccelerator
507 if(!felFieldSetup1_0){
508     static G4LogicalVolume* flogicShape6 = fLXeAccelerator->getlogicShape6();
509     felFieldSetup1_0 = new ElectricFieldSetup();
510     felFieldSetup1_0->SetLocalFieldValue(G4ThreeVector(0.,0.,(4.8*megavolt/m)))
511 ;
511     flogicShape6->SetFieldManager(felFieldSetup1_0->GetLocalFieldManager(),
512     true);
512     G4AutoDelete::Register(felFieldSetup1_0);
513 }
514 if(!felFieldSetup2_0){
515     static G4LogicalVolume* flogicShape10 = fLXeAccelerator->getlogicShape10();
516     felFieldSetup2_0 = new ElectricFieldSetup();
517     felFieldSetup2_0->SetLocalFieldValue(G4ThreeVector(0.,0.,(4.8*megavolt/m)))
518 ;
518     flogicShape10->SetFieldManager(felFieldSetup2_0->GetLocalFieldManager(),
519     true);
519     G4AutoDelete::Register(felFieldSetup2_0);
520 }
521 if(!femFieldSetup1_0){
```

```
522     static G4LogicalVolume* flogicShape1 = fLXeAccelerator->getlogicShape1();
523     femFieldSetup1_0 = new F03FieldSetup();
524     femFieldSetup1_0->SetLocalFieldValue(160.*tesla/(1.*m), G4ThreeVector(),
525     0.*deg);
526     flogicShape1->SetFieldManager(femFieldSetup1_0->GetLocalFieldManager(),
527     true);
528     G4AutoDelete::Register(femFieldSetup1_0);
529 }
530 if(!femFieldSetup2_0){
531     static G4LogicalVolume* flogicShape4 = fLXeAccelerator->getlogicShape4();
532     femFieldSetup2_0 = new F03FieldSetup();
533     femFieldSetup2_0->SetLocalFieldValue(196.*tesla/(1.*m), G4ThreeVector(),
534     90.*deg);
535     flogicShape4->SetFieldManager(femFieldSetup2_0->GetLocalFieldManager(),
536     true);
537     G4AutoDelete::Register(femFieldSetup2_0);
538 }
539 if(!femFieldSetup3_0){
540     static G4LogicalVolume* flogicShape8 = fLXeAccelerator->getlogicShape8();
541     femFieldSetup3_0 = new F03FieldSetup();
542     femFieldSetup3_0->SetLocalFieldValue(179.*tesla/(1.*m), G4ThreeVector(),
543     0.*deg);
544     flogicShape8->SetFieldManager(femFieldSetup3_0->GetLocalFieldManager(),
545     true);
546     G4AutoDelete::Register(femFieldSetup3_0);
547 }
548 if(!femFieldSetup4_0){
549     static G4LogicalVolume* flogicShape12 = fLXeAccelerator->getlogicShape12();
550     femFieldSetup4_0 = new F03FieldSetup();
551     femFieldSetup4_0->SetLocalFieldValue(186.*tesla/(1.*m), G4ThreeVector(),
552     90.*deg);
553     flogicShape12->SetFieldManager(femFieldSetup4_0->GetLocalFieldManager(),
554     true);
555     G4AutoDelete::Register(femFieldSetup4_0);
556 }
557 // LXeModule1
558 if(!felFieldSetup1_1){
559     static G4LogicalVolume* flogicShape6 = fLXeModule1->getlogicShape6();
560     felFieldSetup1_1 = new ElectricFieldSetup();
561     felFieldSetup1_1->SetLocalFieldValue(G4ThreeVector(0.,0.,8.*megavolt/m));
562     flogicShape6->SetFieldManager(felFieldSetup1_1->GetLocalFieldManager(),
563     true);
564     G4AutoDelete::Register(felFieldSetup1_1);
565 }
566 if(!felFieldSetup2_1){
567     static G4LogicalVolume* flogicShape10 = fLXeModule1->getlogicShape10();
568     felFieldSetup2_1 = new ElectricFieldSetup();
569     felFieldSetup2_1->SetLocalFieldValue(G4ThreeVector(0.,0.,8.*megavolt/m));
570     flogicShape10->SetFieldManager(felFieldSetup2_1->GetLocalFieldManager(),
571     true);
572     G4AutoDelete::Register(felFieldSetup2_1);
573 }
```

```
564 }
565 if (!femFieldSetup1_1){
566     static G4LogicalVolume* flogicShape8 = fLXeModule1->getlogicShape8();
567     femFieldSetup1_1 = new F03FieldSetup();
568     femFieldSetup1_1->SetLocalFieldValue(179.*tesla/(1.*m), G4ThreeVector(),
569     0.*deg);
569     flogicShape8->SetFieldManager(femFieldSetup1_1->GetLocalFieldManager(),
570     true);
570     G4AutoDelete::Register(femFieldSetup1_1);
571 }
572 if (!femFieldSetup2_1){
573     static G4LogicalVolume* flogicShape12 = fLXeModule1->getlogicShape12();
574     femFieldSetup2_1 = new F03FieldSetup();
575     femFieldSetup2_1->SetLocalFieldValue(186.*tesla/(1.*m), G4ThreeVector(),
576     90.*deg);
576     flogicShape12->SetFieldManager(femFieldSetup2_1->GetLocalFieldManager(),
577     true);
577     G4AutoDelete::Register(femFieldSetup2_1);
578 }
579
580 // LXeModule2
581 if (!felFieldSetup1_2){
582     static G4LogicalVolume* flogicShape6 = fLXeModule2->getlogicShape6();
583     felFieldSetup1_2 = new ElectricFieldSetup();
584     felFieldSetup1_2->SetLocalFieldValue(G4ThreeVector(0.,0.,11.*megavolt/m));
585     flogicShape6->SetFieldManager(felFieldSetup1_2->GetLocalFieldManager(),
586     true);
586     G4AutoDelete::Register(felFieldSetup1_2);
587 }
588 if (!felFieldSetup2_2){
589     static G4LogicalVolume* flogicShape10 = fLXeModule2->getlogicShape10();
590     felFieldSetup2_2 = new ElectricFieldSetup();
591     felFieldSetup2_2->SetLocalFieldValue(G4ThreeVector(0.,0.,11.*megavolt/m));
592     flogicShape10->SetFieldManager(felFieldSetup2_2->GetLocalFieldManager(),
593     true);
593     G4AutoDelete::Register(felFieldSetup2_2);
594 }
595 if (!femFieldSetup1_2){
596     static G4LogicalVolume* flogicShape8 = fLXeModule2->getlogicShape8();
597     femFieldSetup1_2 = new F03FieldSetup();
598     femFieldSetup1_2->SetLocalFieldValue(179.*tesla/(1.*m), G4ThreeVector(),
599     0.*deg);
599     flogicShape8->SetFieldManager(femFieldSetup1_2->GetLocalFieldManager(),
600     true);
600     G4AutoDelete::Register(femFieldSetup1_2);
601 }
602 if (!femFieldSetup2_2){
603     static G4LogicalVolume* flogicShape12 = fLXeModule2->getlogicShape12();
604     femFieldSetup2_2 = new F03FieldSetup();
605     femFieldSetup2_2->SetLocalFieldValue(186.*tesla/(1.*m), G4ThreeVector(),
606     90.*deg);
606     flogicShape12->SetFieldManager(femFieldSetup2_2->GetLocalFieldManager(),
```



```
    true );
607   G4AutoDelete::Register ( femFieldSetup2_2 );
608 }
609
610 // LXeModule3
611 if (! felFieldSetup1_3){
612   static G4LogicalVolume* flogicShape6 = fLXeModule3->getlogicShape6 ();
613   felFieldSetup1_3 = new ElectricFieldSetup ();
614   felFieldSetup1_3->SetLocalFieldValue (G4ThreeVector (0.,0.,13.4*megavolt/m));
615   flogicShape6->SetFieldManager ( felFieldSetup1_3->GetLocalFieldManager () ,
    true );
616   G4AutoDelete::Register ( felFieldSetup1_3 );
617 }
618 if (! felFieldSetup2_3){
619   static G4LogicalVolume* flogicShape10 = fLXeModule3->getlogicShape10 ();
620   felFieldSetup2_3 = new ElectricFieldSetup ();
621   felFieldSetup2_3->SetLocalFieldValue (G4ThreeVector (0.,0.,13.4*megavolt/m));
622   flogicShape10->SetFieldManager ( felFieldSetup2_3->GetLocalFieldManager () ,
    true );
623   G4AutoDelete::Register ( felFieldSetup2_3 );
624 }
625 if (! femFieldSetup1_3){
626   static G4LogicalVolume* flogicShape8 = fLXeModule3->getlogicShape8 ();
627   femFieldSetup1_3 = new F03FieldSetup ();
628   femFieldSetup1_3->SetLocalFieldValue (179.*tesla / (1.*m) , G4ThreeVector () ,
    0.*deg);
629   flogicShape8->SetFieldManager ( femFieldSetup1_3->GetLocalFieldManager () ,
    true );
630   G4AutoDelete::Register ( femFieldSetup1_3 );
631 }
632 if (! femFieldSetup2_3){
633   static G4LogicalVolume* flogicShape12 = fLXeModule3->getlogicShape12 ();
634   femFieldSetup2_3 = new F03FieldSetup ();
635   femFieldSetup2_3->SetLocalFieldValue (186.*tesla / (1.*m) , G4ThreeVector () ,
    90.*deg);
636   flogicShape12->SetFieldManager ( femFieldSetup2_3->GetLocalFieldManager () ,
    true );
637   G4AutoDelete::Register ( femFieldSetup2_3 );
638 }
639
640 // LXeModuleCCL1
641 if (! felFieldSetup1_4){
642   static G4LogicalVolume* flogicShape1 = fLXeModuleCCL1->getlogicShape1 ();
643   felFieldSetup1_4 = new ElectricFieldSetup ();
644   felFieldSetup1_4->SetLocalFieldValue (G4ThreeVector (0.,0.,12.*megavolt/m));
645   flogicShape1->SetFieldManager ( felFieldSetup1_4->GetLocalFieldManager () ,
    true );
646   G4AutoDelete::Register ( felFieldSetup1_4 );
647 }
648 if (! felFieldSetup2_4){
649   static G4LogicalVolume* flogicShape4 = fLXeModuleCCL1->getlogicShape4 ();
650   felFieldSetup2_4 = new ElectricFieldSetup ();
```

```
651 felFieldSetup2_4->SetLocalFieldValue (G4ThreeVector (0. ,0. ,12.* megavolt/m));
652 flogicShape4->SetFieldManager (felFieldSetup2_4->GetLocalFieldManager () ,
true );
653 G4AutoDelete:: Register (felFieldSetup2_4);
654 }
655 if (! femFieldSetup1_4){
656 static G4LogicalVolume* flogicShape3 = fLXeModuleCCL1->getlogicShape3 ();
657 femFieldSetup1_4 = new F03FieldSetup ();
658 femFieldSetup1_4->SetLocalFieldValue (160.* tesla / (1.*m) , G4ThreeVector () ,
0.* deg);
659 flogicShape3->SetFieldManager (femFieldSetup1_4->GetLocalFieldManager () ,
true );
660 G4AutoDelete:: Register (femFieldSetup1_4);
661 }
662 if (! femFieldSetup2_4){
663 static G4LogicalVolume* flogicShape6 = fLXeModuleCCL1->getlogicShape6 ();
664 femFieldSetup2_4 = new F03FieldSetup ();
665 femFieldSetup2_4->SetLocalFieldValue (160.* tesla / (1.*m) , G4ThreeVector () ,
90.* deg);
666 flogicShape6->SetFieldManager (femFieldSetup2_4->GetLocalFieldManager () ,
true );
667 G4AutoDelete:: Register (femFieldSetup2_4);
668 }
669
670 // LXeModuleCCL2
671 if (! felFieldSetup1_5){
672 static G4LogicalVolume* flogicShape1 = fLXeModuleCCL2->getlogicShape1 ();
673 felFieldSetup1_5 = new ElectricFieldSetup ();
674 felFieldSetup1_5->SetLocalFieldValue (G4ThreeVector (0. ,0. ,12.* megavolt/m));
675 flogicShape1->SetFieldManager (felFieldSetup1_5->GetLocalFieldManager () ,
true );
676 G4AutoDelete:: Register (felFieldSetup1_5);
677 }
678 if (! felFieldSetup2_5){
679 static G4LogicalVolume* flogicShape4 = fLXeModuleCCL2->getlogicShape4 ();
680 felFieldSetup2_5 = new ElectricFieldSetup ();
681 felFieldSetup2_5->SetLocalFieldValue (G4ThreeVector (0. ,0. ,12.* megavolt/m));
682 flogicShape4->SetFieldManager (felFieldSetup2_5->GetLocalFieldManager () ,
true );
683 G4AutoDelete:: Register (felFieldSetup2_5);
684 }
685 if (! femFieldSetup1_5){
686 static G4LogicalVolume* flogicShape3 = fLXeModuleCCL2->getlogicShape3 ();
687 femFieldSetup1_5 = new F03FieldSetup ();
688 femFieldSetup1_5->SetLocalFieldValue (160.* tesla / (1.*m) , G4ThreeVector () ,
0.* deg);
689 flogicShape3->SetFieldManager (femFieldSetup1_5->GetLocalFieldManager () ,
true );
690 G4AutoDelete:: Register (femFieldSetup1_5);
691 }
692 if (! femFieldSetup2_5){
693 static G4LogicalVolume* flogicShape6 = fLXeModuleCCL2->getlogicShape6 ();
```

```
694 femFieldSetup2_5 = new F03FieldSetup();
695 femFieldSetup2_5->SetLocalFieldValue(160.*tesla/(1.*m), G4ThreeVector(),
696 90.*deg);
697 flogicShape6->SetFieldManager(femFieldSetup2_5->GetLocalFieldManager(),
698 true);
699 G4AutoDelete::Register(femFieldSetup2_5);
700 }
701 // LXeModuleCCL3
702 if(!felFieldSetup1_6){
703     static G4LogicalVolume* flogicShape1 = fLXeModuleCCL3->getlogicShape1();
704     felFieldSetup1_6 = new ElectricFieldSetup();
705     felFieldSetup1_6->SetLocalFieldValue(G4ThreeVector(0.,0.,12.*megavolt/m));
706     flogicShape1->SetFieldManager(felFieldSetup1_6->GetLocalFieldManager(),
707 true);
708     G4AutoDelete::Register(felFieldSetup1_6);
709 }
710 if(!felFieldSetup2_6){
711     static G4LogicalVolume* flogicShape4 = fLXeModuleCCL3->getlogicShape4();
712     felFieldSetup2_6 = new ElectricFieldSetup();
713     felFieldSetup2_6->SetLocalFieldValue(G4ThreeVector(0.,0.,12.*megavolt/m));
714     flogicShape4->SetFieldManager(felFieldSetup2_6->GetLocalFieldManager(),
715 true);
716     G4AutoDelete::Register(felFieldSetup2_6);
717 }
718 if(!femFieldSetup1_6){
719     static G4LogicalVolume* flogicShape3 = fLXeModuleCCL3->getlogicShape3();
720     femFieldSetup1_6 = new F03FieldSetup();
721     femFieldSetup1_6->SetLocalFieldValue(160.*tesla/(1.*m), G4ThreeVector(),
722 0.*deg);
723     flogicShape3->SetFieldManager(femFieldSetup1_6->GetLocalFieldManager(),
724 true);
725     G4AutoDelete::Register(femFieldSetup1_6);
726 }
727 if(!femFieldSetup2_6){
728     static G4LogicalVolume* flogicShape6 = fLXeModuleCCL3->getlogicShape6();
729     femFieldSetup2_6 = new F03FieldSetup();
730     femFieldSetup2_6->SetLocalFieldValue(160.*tesla/(1.*m), G4ThreeVector(),
731 90.*deg);
732     flogicShape6->SetFieldManager(femFieldSetup2_6->GetLocalFieldManager(),
733 true);
734     G4AutoDelete::Register(femFieldSetup2_6);
735 }
736 // LXeModuleCCL4
737 if(!felFieldSetup1_7){
738     static G4LogicalVolume* flogicShape1 = fLXeModuleCCL4->getlogicShape1();
739     felFieldSetup1_7 = new ElectricFieldSetup();
740     felFieldSetup1_7->SetLocalFieldValue(G4ThreeVector(0.,0.,12.*megavolt/m));
741     flogicShape1->SetFieldManager(felFieldSetup1_7->GetLocalFieldManager(),
742 true);
743     G4AutoDelete::Register(felFieldSetup1_7);
744 }
```

```
737 }
738 if (!felFieldSetup2_7){
739     static G4LogicalVolume* flogicShape4 = fLXeModuleCCL4->getlogicShape4();
740     felFieldSetup2_7 = new ElectricFieldSetup();
741     felFieldSetup2_7->SetLocalFieldValue(G4ThreeVector(0.,0.,12.*megavolt/m));
742     flogicShape4->SetFieldManager(felFieldSetup2_7->GetLocalFieldManager(),
743     true);
744     G4AutoDelete::Register(felFieldSetup2_7);
745 }
746 if (!femFieldSetup1_7){
747     static G4LogicalVolume* flogicShape3 = fLXeModuleCCL4->getlogicShape3();
748     femFieldSetup1_7 = new F03FieldSetup();
749     femFieldSetup1_7->SetLocalFieldValue(160.*tesla/(1.*m), G4ThreeVector(),
750     0.*deg);
751     flogicShape3->SetFieldManager(femFieldSetup1_7->GetLocalFieldManager(),
752     true);
753     G4AutoDelete::Register(femFieldSetup1_7);
754 }
755 if (!femFieldSetup2_7){
756     static G4LogicalVolume* flogicShape6 = fLXeModuleCCL4->getlogicShape6();
757     femFieldSetup2_7 = new F03FieldSetup();
758     femFieldSetup2_7->SetLocalFieldValue(160.*tesla/(1.*m), G4ThreeVector(),
759     90.*deg);
760     flogicShape6->SetFieldManager(femFieldSetup2_7->GetLocalFieldManager(),
761     true);
762     G4AutoDelete::Register(femFieldSetup2_7);
763 }
764 }
765 // .....oooOO0OOooo.....oooOO0OOooo.....oooOO0OOooo.....oooOO0OOooo
766 .....
767 void LXeDetectorConstruction::SetDimensions(G4ThreeVector dims)
768 {
769     fScint_x = dims[0];
770     fScint_y = dims[1];
771     fScint_z = dims[2];
772     G4RunManager::GetRunManager()->ReinitializeGeometry();
773 }
774 // .....oooOO0OOooo.....oooOO0OOooo.....oooOO0OOooo.....oooOO0OOooo
775 .....
776 void LXeDetectorConstruction::SetHousingThickness(G4double d_mtl)
777 {
778     fD_mtl = d_mtl;
779     G4RunManager::GetRunManager()->ReinitializeGeometry();
780 }
781 // .....oooOO0OOooo.....oooOO0OOooo.....oooOO0OOooo.....oooOO0OOooo
782 .....
783 void LXeDetectorConstruction::SetNX(G4int nx)
```

## Section

```
781 {
782     fNx = nx;
783     G4RunManager::GetRunManager()->ReinitializeGeometry();
784 }
785
786 // .....
787
788 void LXeDetectorConstruction::SetNY(G4int ny)
789 {
790     fNy = ny;
791     G4RunManager::GetRunManager()->ReinitializeGeometry();
792 }
793
794 // .....
795
796 void LXeDetectorConstruction::SetNZ(G4int nz)
797 {
798     fNz = nz;
799     G4RunManager::GetRunManager()->ReinitializeGeometry();
800 }
801
802 // .....
803
804 void LXeDetectorConstruction::SetPMTRadius(G4double outerRadius_pmt)
805 {
806     fOuterRadius_pmt = outerRadius_pmt;
807     G4RunManager::GetRunManager()->ReinitializeGeometry();
808 }
809
810 // .....
811
812 void LXeDetectorConstruction::SetDefaults()
813 {
814     // Resets to default values
815     fD_mtl = 0.0635 * cm;
816     fD_moderator = 3. * cm;
817
818     fScint_x = 6. * cm;
819     fScint_y = 6. * cm;
820     fScint_z = 20. * cm;
821
822     fNx = 1;
823     fNy = 1;
824     fNz = 0;
825
826     fOuterRadius_pmt = 2.3 * cm;
827
828     fSphereOn = false;
```

```
829   fRefl      = 1.0;
830
831   fNfibers   = 1;
832   fWLSslab   = false;
833   fMainVolumeOn = true;
834   fMainVolume = nullptr;
835   fSlab_z     = 2.5 * mm;
836
837   G4UImanager::GetUIpointer()->ApplyCommand(
838     "/LXe/detector/scintYieldFactor 1.");
839
840   if (fLXe_mt)
841     fLXe_mt->AddConstProperty("SCINTILLATIONYIELD", 12000. / MeV);
842   if (fMPTPStyrene)
843     fMPTPStyrene->AddConstProperty("SCINTILLATIONYIELD", 10. / keV);
844 }
845
846 // .....oooOO0OOooo ..... oooOO0OOooo ..... oooOO0OOooo ..... oooOO0OOooo
847     .....
848 void LXeDetectorConstruction::SetSphereOn(G4bool b)
849 {
850   fSphereOn = b;
851   G4RunManager::GetRunManager()->ReinitializeGeometry();
852 }
853
854 // .....oooOO0OOooo ..... oooOO0OOooo ..... oooOO0OOooo ..... oooOO0OOooo
855     .....
856 void LXeDetectorConstruction::SetHousingReflectivity(G4double r)
857 {
858   fRefl = r;
859   G4RunManager::GetRunManager()->ReinitializeGeometry();
860 }
861
862 // .....oooOO0OOooo ..... oooOO0OOooo ..... oooOO0OOooo ..... oooOO0OOooo
863     .....
864 void LXeDetectorConstruction::SetWLSslabOn(G4bool b)
865 {
866   fWLSslab = b;
867   G4RunManager::GetRunManager()->ReinitializeGeometry();
868 }
869
870 // .....oooOO0OOooo ..... oooOO0OOooo ..... oooOO0OOooo ..... oooOO0OOooo
871     .....
872 void LXeDetectorConstruction::SetMainVolumeOn(G4bool b)
873 {
874   fMainVolumeOn = b;
875   G4RunManager::GetRunManager()->ReinitializeGeometry();
876 }
```

## Section

```
877
878 // .....oooOO00O0ooo ..... oooOO00O0ooo ..... oooOO00O0ooo ..... oooOO00O0ooo
      .....
879
880 void LXeDetectorConstruction::SetNFibers(G4int n)
881 {
882     fNfibers = n;
883     G4RunManager::GetRunManager()->ReinitializeGeometry();
884 }
885
886 // .....oooOO00O0ooo ..... oooOO00O0ooo ..... oooOO00O0ooo ..... oooOO00O0ooo
      .....
887
888 void LXeDetectorConstruction::SetMainScintYield(G4double y)
889 {
890     fLXe_mt->AddConstProperty("SCINTILLATIONYIELD", y / MeV);
891 }
892
893 // .....oooOO00O0ooo ..... oooOO00O0ooo ..... oooOO00O0ooo ..... oooOO00O0ooo
      .....
894
895 void LXeDetectorConstruction::SetWLSScintYield(G4double y)
896 {
897     fMPTPStyrene->AddConstProperty("SCINTILLATIONYIELD", y / MeV);
898 }
899
900 // .....oooOO00O0ooo ..... oooOO00O0ooo ..... oooOO00O0ooo ..... oooOO00O0ooo
      .....
901
902 void LXeDetectorConstruction::SetSaveThreshold(G4int save)
903 {
904     // Sets the save threshold for the random number seed. If the number of
905     // photons generated in an event is lower than this, then save the seed for
906     // this event in a file called run###evt###.mdm
907
908     fSaveThreshold = save;
909     G4RunManager::GetRunManager()->SetRandomNumberStore(true);
910 }
911
912 // .....oooOO00O0ooo ..... oooOO00O0ooo ..... oooOO00O0ooo ..... oooOO00O0ooo
      .....
```

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- [1] World Health Organisation
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