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TRITIUM: Design, construction and commissioning of an in-water tritium detector

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*Dedicated to
my family*

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Ver memoria de "Detectores monolíticos y sensores compatibles con altos campos magnéticos para tomografía por emisión de positrones"

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IV

Gente de ATLAS, URMILA, CARLOS MARÍÑAS

Y PENSAR GENTE QUE ME DEJO POR EL CAMINO.

Abstract

Tritium is one of the most frequently emitted radioisotopes in a nuclear power plant. Large quantities of tritium are normally produced in the water of their cooling system, which are finally emitted to the environment. Due to the fact that high quantities of tritium could be dangerous for human health and for the environment, there exist several legislations around the world which try to control this radioactive emissions in each country, like the Directive European 2013/51/Euratom, which establishes the tritium limit in drinking water in Europe at 100 Bq/L, or the U. S. Environmental Protection Agency, in United States, whose tritium limit in drinking water is established at 740 Bq/L.

Nowadays, due to such a low energy emitted in the tritium decay, we need high sensitive detectors for measuring it like LSC. The problem with LSC is that it is a off-line method whose measurement process can take up to 3 or 4 days, too much time if there are any problem with the NPP.

Detectors based on solid scintillators is a promissing idea for building a tritium detector that works in quasi-real time. This type of detectors has been developed so far succesfully but without achieving enough sensibility for measuring the legal limits.

In this study the results of TRITIUM project is presented. In the

framework of this project we have developed a quasi-real time monitor for low tritium activities in water. This monitor is based on a tritium detector that contains several detection cells which we read in parallel, several active vetos and a pasive shielding for reduce the natural background of our system and an ultra pure water system to prepare the sample before we measure. Each detection cell is made up of hundreds of scintillating fibers read out by PMTs or SiPM arrays.

The final objective of this monitor will be the radiological protection around the nuclear power plant. This monitor will provide an alarm in case of an unexpected tritium release. It will be included in the early alarm system of Extremadura consisting of several detectors whose objective is to reduce the impact of Nuclear Power Plants to the environment.

Keywords: una, dos, tres...

Nomenclature and Acronyms

Acronyms:

| | | |
|---------------------|---|---|
| <i>OECD</i> | — | Organisation for Economic Co-operation and Development |
| <i>UN</i> | — | United Nations |
| <i>UNFCCC</i> | — | The United Nations in the framework convention about Climate change |
| <i>ITER</i> | — | International Thermonuclear Experimental Reactor |
| <i>NPP</i> | — | Nuclear Power Plants |
| <i>DOE</i> | — | Department of Energy |
| <i>U.S.A.</i> | — | United States of America |
| <i>U.S.</i> | — | United States |
| <i>EPA</i> | — | Environmental Protection Agency |
| <i>LDL</i> | — | Lower Detection Limit |
| <i>PWR</i> | — | Pressurized Water Reactor |
| <i>BWR</i> | — | Boiled Water Reactor |
| <i>HWR</i> | — | Heavy Water Reactor |
| <i>GCR</i> | — | Gas-Cooled Reactor |
| <i>PHWR</i> | — | Pressurized Heavy Water Reactor |
| <i>quasi – real</i> | — | Less than 10 minuts |
| <i>LSC</i> | — | Liquid Scintillation Counting |
| <i>LWR</i> | — | Liquid Water Reactor |
| <i>IC</i> | — | Ionization chamber |
| <i>BIXS</i> | — | Beta Induced X-ray Spectrometry |

VIII

| | | |
|--------------|---|---|
| <i>PMT</i> | — | Photoelectron Multiplier Tube |
| <i>SDD</i> | — | Silicon Drift Detector |
| <i>EEC</i> | — | European Economical Community |
| <i>SiPM</i> | — | Silicon Photoelectron Multiplier |
| <i>CNRS</i> | — | Le Centre National de la Recherche Scientifique, France |
| <i>ALARA</i> | — | As Low As Reasonably Achievable |
| <i>CCD</i> | — | Charge-Coupled device |
| <i>HV</i> | — | High Voltage |
| <i>QE</i> | — | Quantum Efficiency |
| <i>CE</i> | — | Collection Efficiency |

Atomic and nuclear symbols

| | | |
|------------------------|---|---|
| CO_2 | — | Carbon Dioxide |
| CH_4 | — | Methane |
| N_2O | — | Nitrous oxide |
| HFC | — | Hydrofluorocarbons |
| PFC | — | Perfluorocarbons |
| SF_6 | — | Sulfur Hexafluoride |
| ^1_1H | — | Hydrogen |
| ^2_1H | — | Deuterium (Non-radioactive hydrogen isotope, 1 neutron) |
| ^3_1H | — | Tritium (radioactive hydrogen isotope, 2 neutrons) |
| ^4_2He | — | Helium |
| ^3_2He | — | Isotope of the Helium(Non-radioactive, 1 neutrons) |
| $^{14}_7\text{N}$ | — | Nitrogen |
| $^{12}_6\text{C}$ | — | Carbon |
| ^6_3Li | — | Lithium isotope |
| ^7_3Li | — | Lithium |
| $^{10}_5\text{B}$ | — | Boron |
| $^{16}_8\text{O}$ | — | Oxygen |
| $^{222}_{86}\text{Rn}$ | — | Radon |

| | | |
|------------------------|---|--|
| $^{40}_{19}\text{K}$ | — | Potassium |
| $^{137}_{55}\text{Cs}$ | — | Cesium |
| n | — | free neutron |
| H_2O | — | Usual water |
| D_2O | — | Heavy water |
| HDO | — | Semi heavy water |
| $T_{1/2}$ | — | Half-life time of a radioactive element |
| β | — | Beta decay |
| $\bar{\nu}_e$ | — | Electron antineutrino |
| e^- | — | Electron |
| γ | — | Gamma |
| p | — | Proton |
| α | — | Alpha (nucleus with two protons) |
| σ_i | — | Cross section of the "i" process |
| η_{det} | — | Intrinsic detector efficiency |
| F_{sci} | — | Active surface of the plastic scintillator |
| ε_{det} | — | Specific detector efficiency |
| mip | — | minimum ionizing particle |

Units:

| | | |
|----------------------------|---|---|
| Btu | — | British thermal unit |
| STP | — | Standard temperature (0°C) and pressure (1 atm) |
| W | — | watt |
| h | — | hour |
| $\text{g CO}_2/\text{kWh}$ | — | grams of CO_2 per kilowatt hour |
| L | — | Liter |
| Bq | — | Becquerel, Nuclear decay number per second |
| Bq/L | — | Becquerel per liter |
| Ci | — | Curios |
| Ci/L | — | Curios por litro |

| | | |
|------------|---|----------------------------|
| yr | — | year |
| $\mu S/cm$ | — | MicroSivers per centimeter |

Añadir en un futuro:

| | | |
|--------|---|--------------------------------------|
| $D\&D$ | — | Decontamination and Decommissioning. |
| DWS | — | Drinking water standars |
| UDL | — | Upper Detection Limit |
| NA | — | Numerical Apertures |
| $PMMA$ | — | Polymethyl Methacrylates |
| UV | — | Ultraviolet |
| WLS | — | Wavelength shifter |
| T | — | Temperature ($^{\circ}C$). |
| V | — | Volume (m^3). |

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Chapter 1

Introduction

1.1 Global energy context

The energy necessities around the world has been increased a 60% in the last 25 years and they are growing each day due to, mainly, the strong population growth (mundial population has been increased in a 40% between 1990 and 2015) and the fast development of emerging countries like China, India and Brazil. [1].

On top of that the outlook is that these energy necessities will keep increasing as you can see in the figure 1.1a, specially for countries which don't belong to the OECD like countries which I have said before. In this figure you can see that the energy quantity which will be used on 2040 is expected to be the double of the one which we used in 2000.

Nowadays, as you can see in the figure 1.1b, the most used elements for getting the energy that we use are liquid fuels, coal and natural gas, namely, natural elements. This fact has two problems. On the one hand, natural elements are limited and this is a problem due to the huge

increasing of the energy consumed in the world and, on the other hand, obtaining energy from these natural elements produces large amounts of greenhouses gases (mainly CO_2) which contribute to the environmental contamination, global warming, deforestation, etc. and this is a very big problem, specially right now, since United Nations (UN) did a communication on November 25, 2019 [3], where they claimed that the lifestyle of the world population has not changed in the last years and, as a reason of that, we achieved the highest level of the CO_2 in the history.

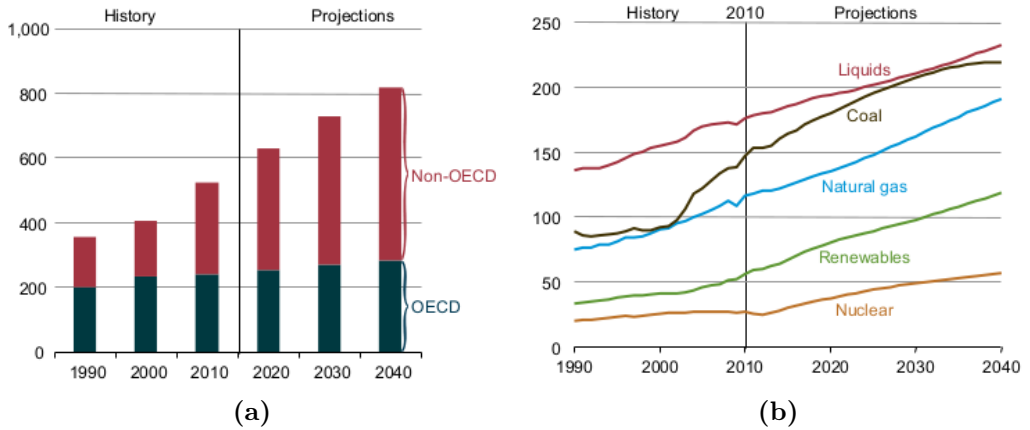


Figure 1.1 – World energy consumption from 1990 up to now and outlooks for the future until 2040 (units in 10^{15} Btu) [2]

In order to control these emissions, the United Nations in the Framework Convention about Climate Change (UNFCCC), has developed a protocol, whose name is Kyoto [4]. The objective of this protocol is to control and reduce the global negative environmental impact. It is focused on 6 different gases, carbon dioxide (CO_2), methane (CH_4), nitrous oxide (N_2O) and other three types of fluorinate industrial gases (HFCs, PFCs and SF_6), which are related with the greenhouse effect and whose consequence is the global warming. This convention only encourage to belonging countries to the United Nations to reduce their greenhouses gases emissions but this protocol commits the countries who has signed it to do so.

Therefore, we currently have a problem because, on the one hand, we want to maintain, even increase, this economic growth and for that we need to produce as much energy as we can but, on the other hand, we need to reduce the negative environmental impact. Hence, what we need is a energy source with which we can get a big quantity of energy with a very low greenhouse gases emissions. One possibility which we have is the nuclear fusion plants. They don't emit greenhouse gases and with its energy, which is practically limitless, we could satisfy the humanity's energy necessities .

The largest representative in this sector is the ITER [5] (International Thermonuclear Experimental Reactor) that is going to be the largest fusion reactor in the world. The ITER is currently under construction in Cadarache, France, and its objective is to provide the concept of sustained fusion. The nuclear reaction which ITER try to reproduce in the earth is the one that uses deuterium (${}^2_1\text{H}$) and/or tritium (${}^3_1\text{H}$) because it is the only one whose requirements of temperature (several hundreds of millions degrees) and pressure we can fulfill in the earth. Among the possible combinations between both (table 1.1), the choice of ITER is the nuclear reaction between deuterium and tritium in a 50:50 mixture because, as you can see in this table, it is the nuclear reaction with which we get the highest energy release.

| Reaction | Products | Energy gain (MeV) |
|-------------------------------|-------------------------------|-------------------|
| ${}^2\text{H} + {}^3\text{H}$ | ${}^4\text{He} + \text{n}$ | 17.6 |
| ${}^3\text{H} + {}^3\text{H}$ | ${}^4\text{He} + 2\text{n}$ | 11.3 |
| ${}^2\text{H} + {}^2\text{H}$ | ${}^3\text{H} + {}^1\text{H}$ | 3.98 |
| ${}^2\text{H} + {}^2\text{H}$ | ${}^3\text{He} + \text{n}$ | 3.25 |

Table 1.1: Fusion reactions between deuterium and tritium[6]

Nevertheles, the fusion power plants is not already prepared for working, they are still in experimental phase and its researchers need to solve some important problems like inestabilitiy vortex, materials which

withstand such high temperatures, etc. [7]. Hence, although we know that the nuclear fusion plants will be the energy of the future, nowadays, we have to wait until all their problems has been resolved.

Other possible energy source is the existing Nuclear Power Plants (NPPs). With the NPPs we can practically avoid the problem of the greenhouse gases emission. We have to take into account that, although the nuclear fission reaction doesn't emit greenhouse gasses, the total proces to obtain the energy, which involves the uranium mining and milling, transport, uranium enrichment, etc., has a small contribution to the annual release of greenhouses gases. These emissions are difficult to estimate because, on the one hand, they depend on the NPP that we consider (for instance, there are studies which show that asian NPPs has higher emissions [8]) and, on the other hand, there are some tasks whose greenhouse gases emission are difficult to quantify[8].

There's exist a study [8] which analyzes 19 different studies of different NPPs. His estimation for the total greenhouses gases emission of a NPP is 66 g CO₂/kWh which was obtained as a average of these 19 studies considered. In the table 1.2 this estimation is compared with the estimation for other energy kinds. There you can check that, the emissions due to NPPs is much more smaller, one order or more, than the emissions from burning natural elements.

NPPs are already working and, nowadays, they are essential for providing a big part of the electric power that is used in the current world (more than a 20% in Spain as you can see in the table 1.3).

NPP is one of the cheapest source. It is a stable source which doesn't depend on meteorological parameters and, although there are other alternative energy sources which are being developed quickly (photovoltaic, wind, tidal energy, etc.), even other concepts of energy production and saving (local production, solar roofs, energy efficiency, smart cities, etc.),

| Technology | Estimate (g CO_2 /kWh) |
|---------------|--------------------------|
| Wind | 9 – 10 |
| Hydroelectric | 10 – 13 |
| Biogas | 11 |
| Solar thermal | 13 |
| Biomass | 14 – 41 |
| Solar PV | 32 |
| Geothermal | 38 |
| Nuclear | 66 |
| Natural gas | 443 |
| Fuel cell | 664 |
| Diesel | 778 |
| Heavy oil | 778 |
| Coal | 960 – 1050 |

Table 1.2: Estimations of CO_2 emissions for several kinds of energy sources[8]

| Type of energy source | Contr. | Type of energy source | Contr. |
|-----------------------|--------|--|--------|
| Nuclear | 22.0% | Wind | 20.9% |
| Coal | 4.2% | Hydraulics | 9.7% |
| Combined Cycle | 20.1% | Solar Photovoltaic | 3.5% |
| Cogeneration | 11.8% | Solar thermal | 2.0% |
| No-renewable waste | 0.8% | Other renewables | 1.4% |
| Pumping turbine | 0.6% | renewable waste | 0.3% |
| | | Imputed balance of international exchanges | 2.7% |

Table 1.3: Contribution of each energy source to the total energy consumed in Spain in 2019 [9]

today they are not develop enough to fully supply ourselves with them.

The detractors of nuclear energy argue that NPPs facilitate nuclear proliferation or there are a risk of radioactive contamination and accidents like it happened in the past: Chernobyl, Fukushima and other accidents with lesser impact such as Three Mile Island, near to Pensilvania, USA [10].

Although we know that the nuclear energy is not the energy of the future since it produces nuclear waste which, by the moment, we don't know how we can eliminate, it is difficult that we leave to use nuclear energy because, now, we don't have a better solution for obtaining the energy which we need.

In Spain the government are doing an effort in order to remove all nuclear power plants since they are not going to build new nuclear reactors and they are only waiting until the NPPs have reached the end of their useful life (approximately 40 years). They expect to close all NPPs between 2020 and 2030 [11].

Other countries like France, where the 77% of their energy consumed is obtained from nuclear sources, prefer to maintain their nuclear facilities and there's even exist other countries that believe that nuclear energy is a safe investment like China who announced in 2016 that they were going to build 60 new nuclear reactors in the next dedade [12] or USA, who made an investment of 35 million of euros in 2019 for development and improvement of nuclear power plants [13].

In any case it is not important if we agree or not with nuclear energy source. The only important thing is that the nuclear energy production in the world is not going to stop in the next decade, in fact, it will increase as you have seen in the outlooks of the figure 1.1b. Therefore the development of different types of alarm systems is a good investment of both, time and money. Safety is not a negotiable aspect and there must

be mechanisms that warn us of any malfunction of a nuclear power plant. Hence, our work has based on the development of a monitor that we can use as early alarm in case of any problem happen in a NPP.

Generally, a nuclear reactor, which is working in normal mode, is characterized by extreme stability and, therefore, by a constant emission of radioactive isotopes so the first alarm signs of any malfunctioning of a NPP is a variation of this radioactive emission rate.

Between all the radioactive elements which is produced in a nuclear power plant, the most frequently produced is tritium as DOE complex [14] [15] and other research facilities in China [16] have seen in their installations and in ground water, surface water, and process waste water around their facilities. However, as we will see in the section 1.3, the current methods which is used for monitoring this radioactive element has some limitations.

For these reasons, the radioactive element which we have chosen for monitoring with our early alarm system is tritium. We have focused our alarm system for working with NPPs but it could be also interesting for other tasks where tritium is involved like monitoring the behaviour of future fusion nuclear plants (ITER will need up to several tens of kilograms of tritium for working, which correspond to several TBq of tritium) or any nuclear research facility (tritium is a common emission of these places [17],[18]), tracking the movement of tritium contaminated plumes in ground water [19] or demonstrate the compliance with the government agencies which fix the limit of the emitted radionuclides to the environmental.

We have to take into account that the limit of the emission of each radioactive element depends on the government agency who manages it, and the regulation directives that is implemented in that place so, as a consequence, it is different in each country. For example, in Europe, the agency that manages these limits is Council Directive and the limit which they have established for tritium in drinking water is $A = 100 \text{ Bq/L}$ [20]. In

USA, the organization is United States Environmental Protection Agency (U. S. EPA) and this limit is $A = 20 \text{ nCi/L} = 740 \text{ Bq/L}$ [21].

Tritium is normally produced in the water that there are in the cooling system or the moderator of some NPPs. It usually appear by neutron capture of the deuterium, which exist in the heavy water (D_2O), semi-heavy water (HDO) or the deuterium which has created by neutron capture in usual water (H_2O). All these processes have a big probability to happen due to the huge quantity of neutrons which we has in the nuclear reactor, $10^{14} \text{ n cm}^{-2}\text{s}^{-1}$ [22].

The quantity of tritium will be different for each reactor type because, as we will see in the next seccion, the cross section of tritium production will depend on the materials which there are in any type of NPP. In the table 1.4 we can see the emissions of different types of nuclear reactors:

| Reactor type | Gaseous discharge (GBq/y) | Liquid discharge (GBq/y) |
|--------------|---------------------------|--------------------------|
| PWR | $3.70 \cdot 10^3$ | $2.59 \cdot 10^4$ |
| BWR | $1.85 \cdot 10^3$ | $3.70 \cdot 10^3$ |
| HWR | $7.40 \cdot 10^5$ | $1.85 \cdot 10^5$ |
| GCR | $7.40 \cdot 10^3$ | $1.11 \cdot 10^4$ |

Table 1.4: Emission of tritium in different types of nuclear reactors[16]

The tritium which is created in the water of the cooling system is finally released partially or totally to the environment. Between these types, the most common way is HTO [16].

Our alarm system will monitor the activity of tritium in the water of the cooling system of the NPP which is emitted to the environmental. It can also work for the water of the moderator in these types of NPP but it's not our objective because it's a close circuit so it's not a emission (unless the moderator is leaking, in which case our alarm system would detect it indirectly due to a variation of tritium activity in the water of the cooling

system released to the environment).

The measurement of the tritium activity is one of the systematic environmental control which is performed during energy production by NPPs. It is normally done by LSC technic which has a very good detection capability and precision but it has some problems, for example, it needs too long time for taking a measurement (2 days or more). I will speak more about LSC technic in the section 1.3.

The detection of this tritium in quasi-real time (< 10 min) is important because of the following reasons:

1. It can warn us about the production of an excessive number of neutrons in the nuclear reactor due to the overheating of itself or a leakage of the water from the primary circuit of the cooling system in a nuclear power plant due to some break (perhaps because of an excessive preassure, other alarm sign of a malfunctioning of a nuclear reactor). Both causes could become in a very dangerous problems so the tritium detection in quasi-real time could be important in order to quickly detect and to solve it.
2. The water, which there are in the secondary circuit of the cooling system, will be released to the enviromental, usually rivers or seas, after using it for cooling purposes. Generally this water will be used later for human consuming, irrigation of all kind of plantations or it will arrive to a places where we fish.

Due to such a low legal limit in comparation with the activities of tritium inside of a nuclear reactor, it is possible that, if the nuclear power plant don't work correctly, the activity of tritium water released exceeds this limit and it become this water in no drinkable and these crops into inedibles. On top of that, the life time of tritium is more than 12 years so these places will remain contaminated during a lot of time.

3. There's exist a lot of rivers or seas, which is used for cooling systems of the nuclear power plants, that are crossborders, that's, they are shared by several countries like our case as we will see in the seccion 1.4. The emisión of an excessive tritium activity of one country could be affect severely to the other country creating new international conflicts between them.

Because of all these reasons it is very important that we have an alarm system which is capable of measuring such low tritium activities in quasi-real time. Nevertheless, as we will see in the section 1.3, currently there are not any technic with which we can fulfill these requeriments.

All these reasons has motivated the *Tritium* project, whose objective is the development of a system for quasi-real time monitoring of low radioactive levels of tritium in water for security applications in nuclear power plants.

1.2 Tritium properties

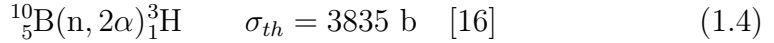
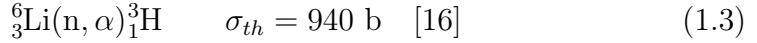
Tritium is the only radioactive isotope of hydrogen. It was first time produced in 1934 from neutron capture of deuterium by Ernest Rutherford, Mark Oliphant and Paul Harteck [23] and it was first time isolated in 1939 by Luis Walter Alvarez and Robert Cornog [24], who checked that tritium is a radioactive element.

Tritium can be found in the environment since it is normally produced through the interaction of cosmic rays and gaseous elements of the upper atmosphere like nitrogen ($^{14}\text{N}(\text{n}, ^3\text{H})^{12}\text{C}$) [25] and oxygen ($^{16}\text{O}(\text{n}, ^3\text{H})^{14}\text{N}$) [26]. Then tritium becomes water (HTO) and reaches the earth's surface as rain with an estimated production rate of $4 \cdot 10^6$ Ci/yr ($1.48 \cdot 10^8$ GBq/yr) [16] [25] .

Tritium can be produced artificially in the environment from many different anthropogenic origins. There are a big amount of tritium which was produced on militar nuclear test explosions between 1945 and 1975, whose estimated production rate is $8 \cdot 10^9$ Ci ($2.96 \cdot 10^{11}$ GBq) and a part of that still remain. It was mainly produced from the nuclear reactions $^{14}\text{N}(\text{n}, ^3\text{H})^{12}\text{C}$ and $^2\text{H}(\text{n}, \gamma)^3\text{H}$. Tritium can be also produced by commercial producers of radiolumincent and neutron generator devices ($1 \cdot 10^6$ Ci/yr), nuclear power and defense industries (less than $2 \cdot 10^6$ Ci/yr), several research facilities and nuclear reactor operation ($2 \cdot 10^6$ Ci/GWyr), whose main production channels are [16] [25]:

$$^2_1\text{H}(\text{n}, \gamma)^3_1\text{H} \quad \sigma_{th} = 5.2 \cdot 10^{-4} \text{ b} \quad [16] \quad (1.1)$$

$$^3_2\text{He}(\text{n}, \text{p})^3_1\text{H} \quad \sigma_{th} = 5330 \text{ b} \quad [16] \quad (1.2)$$



Tritium is a radioactive element whose half-life time is $T_{1/2} = 12.32$ years. It has one proton and two neutrons and decays exclusively through β radiation, that's, it doesn't have other type of radioactive decay. In this decay, one neutron of tritium is transformed in a proton plus electron and electron-antineutrino according to the following equation:



Then the nucleus of the tritium son has two protons and one neutron so it is a helium isotope, ${}^3_2\text{He}$ which is stable. Therefore, the nuclear reaction which descript the β^- decay of the tritium is:



In the Figure 1.2 we can see the scheme of tritium energy levels. In this decay we practically don't have the possibility of detecting the neutrinos because it interacts very weakly with matter and, therefore, with our detector ($\sigma \propto 10^{-42} \text{ cm}^2$ [27]) and, since ${}^3\text{He}$ has a much larger mass than electrons and neutrinos, by conservation of energy and momentum, the energy that is taken by its atom is very small. Therefore, we will focus on the electron detection.

The energy that is released in this nuclear reaccion is constant, 18.6 keV, but it is divided between the products of this reaccion. Therefore

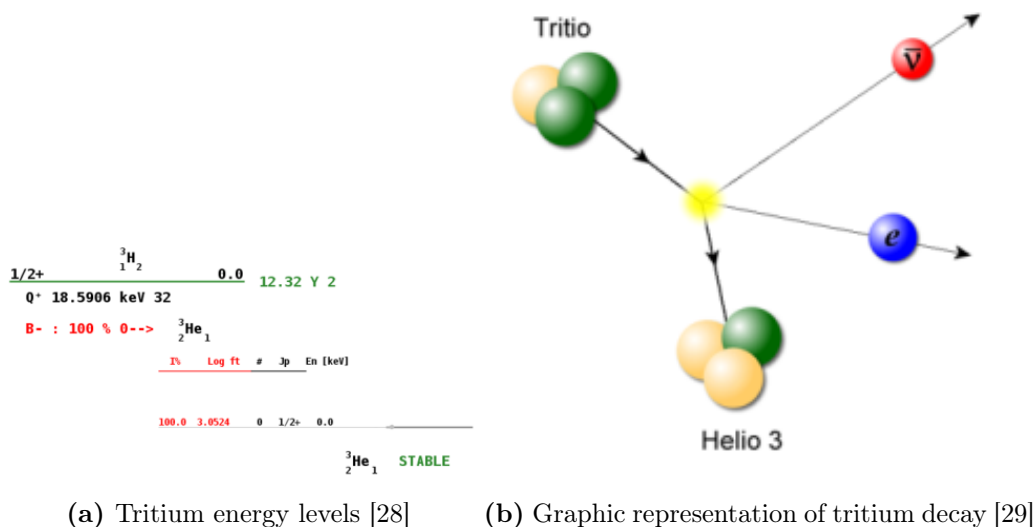


Figure 1.2 – Tritium decay

not all beta particle (electrons) will have the maximum energy. This is what we can see in the figure 1.3, which is the energetic spectrum of the electrons which are emitted in the tritium decay. The maximum energy of this electrons is 18.6 keV (when beta particles have all the energy), which is the endpoint energy of this spectrum, the average energy is 5.7 keV and the most likely value is slightly below of the average energy, around 4.5 keV.

Keep in mind that, although the helium isotope is stable, it will be exited immediately after this decay. As a consequence, after the tritium β^- decay, we will have a subsequent dexcitation of the ${}^3\text{He}$ which will produce fotons, γ , with several well-defined energies that correspond to their energy levels, X-rays. COMPROBAR... It doesn't affect directly to our detector due to the efficiency of our detector at those wavelengths, as we will se in the chapter ??, section ??, but it could be affect indirectly. BUSCAR ESTAS ENERGÍAS Y VER SI SON TAN DIFERENTES COMO PARA NO AFECTAR DIRECTAMENTE O SE PARENCE A LAS DEL TRITIO Y SI AFECTAN DIRECTAMENTE.

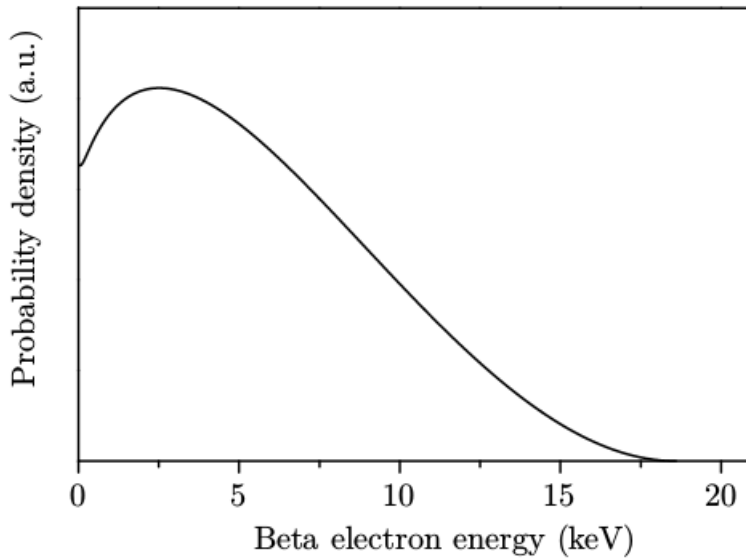


Figure 1.3 – Energy spectrum of tritium electrons [30]

The releasing energy, which is produced in the tritium decay, is very little. In fact, it is the radioactive isotope with the lowest energy released in its β disintegration [25]. As a consequence the β particles which is emitted in this tritium decay will have a very little mean free path as you can see in the table 1.5.

On the one hand, it means that the tritium electrons is easily stopped even for simply walls like our clothes, the laboratories gloves or even the our skin it-self, that's, the radioactive hazard is low. Nevertheless, the danger of tritium is increased when tritium is ingested or inhaled because if it has enough radioactivity it can affect to our internal organs because it has a high biologic life time, 9.5 days [25], time during which tritium remains in our body and we will be receiving radioactive dose due to tritium radiation. Therefore, their health hazard is high.

On the other hand, this short mean free path will be a problem when we try to detect tritium and due to that, there are some limitations

| Material | Energy (β)(keV) | Penetration Depth |
|--|-------------------------|--------------------|
| ${}^3_1\text{H}_2$, STP | 5.7 | 0.26 cm |
| ${}^3_1\text{H}_2$, STP | 18.6 | 3.2 cm |
| Air, STP | 5.7 | 0.036 cm |
| Air, STP | 18.6 | 0.45 cm |
| Water, soft tissue (solid matter whose density is $1 \text{ g} \cdot \text{cm}^{-3}$) | 5.7 | $0.42 \mu\text{m}$ |
| Water, soft tissue (solid matter whose density is $1 \text{ g} \cdot \text{cm}^{-3}$) | 18.6 | $5.2 \mu\text{m}$ |

Table 1.5: Mean Free Path of tritium isotope for several energies and several environments [25]

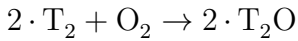
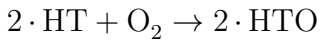
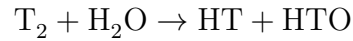
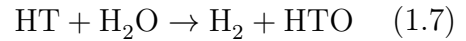
which we will have to take into account when we design our detector.

Tritium has different physical properties than other natural isotopes of the hydrogen like different boiling points as you can see in the table 1.6 or the property of auto-radiolysis which only happens when radioactive elements are presented. The auto-radiolysis exists because the energy released in tritium decay is larger than the energy bond of oxygen and hydrogen in water molecules (5.2 eV) or the ionization energy of water molecules (12.6 eV) so it can break up these molecules [31]. Due to the auto-radiolysis, some radicals appear in the water whose corrosivity is increased and it is something that we have to take into account when choosing the materials that will make up our detector.

Although tritium has different physical properties it has almost the same chemical behaviour than other hydrogen isotopes. Tritium, like hydrogen, is a gas at standard conditions of temperature (273 K) and pressure (1 atm) forming a two-atom molecules which can be HT, DT and T₂. It can become in tritium water through oxidation and exchange reactions as you can see in the following chemical equations[25]:

| Molecule | Boiling point (for gases) (K) | oxidation form |
|----------------|-------------------------------|------------------|
| H ₂ | 20.39 | H ₂ O |
| HD | 22.14 | HDO |
| HT | 22.92 | HTO |
| D ₂ | 23.66 | D ₂ O |
| DT | 24.38 | DTO |
| T ₂ | 25.04 | T ₂ O |

Table 1.6: Gas molecules of hydrogen isotopes and their boiling points

Oxidation :*Exchange*

Due to this chemical similarity tritium water can perform the same chemical process than non-radioactive water, some times with higher rate if the tritium concentration is high enough to catalyze the reaction. Its biological hazard comes from this chemical similarity since tritium water is able to substitute normal water in human body. On top of that, tritium water has a higher absorption in human body, around 99%, than tritium gas, whose absorption in the human body is less than $5 \cdot 10^{-3}\%$ by inhalation or practically negligible by skin absorption [25] so it is more dangerous.

1.3 State-of-the-art in tritium detection

Measurement of tritium activity is one of the systematic environmental controls that have been carried out for dozens of years around nuclear power plants during their energy production and around nuclear research facilities.

As a consequence, this measurement has been attempted with many different technologies so far in order to improve the state of the art of each time. The most researched techniques are summarized in the table 1.7.

| | LSC | IC | Calorimetry | BIXS |
|-------------------|-----------------------|--------------------|-------------|------------|
| Measured quantity | Scintillation photons | Ionization current | heat | X-rays |
| LDL | \sim Bq | 10 – 100 kBq | \sim GBq | \sim MBq |
| Sample form | Liquid | Gas, vapor | All | All |

Table 1.7: State-of-the-art in the tritium detection for different technics [30]

Nowadays, the most used technic for measuring tritium in water is the LSC. It consists of mixing a liquid sample (some ml for environmental measurements or less for higher activities) with liquid scintillator. In our laboratories, LARAM, at the University of Valencia, this mixture is made in a ratio of 50:50 [32] but it will depend on each system and each sample [33] [34]. In this technic, the β energy that is emitted from the sample excites the molecular energy levels of the liquid scintillator and it is quickly desexcited emitting several photons with a well-know energy (fluorescence), normally in the visible range. Finally these photons are detected with photosensors, processed and analysed.

This technic has a very good detection capability and precision (LDL for tritium better than 1 Bq/L [35]) but it has some problems. On the one hand it need too time for taking a mesurment (more than 2 days) and,

on the other hand, although this sample could be non-radioactive, it contain toluene which is a toxical chemical waste so we need to follow a special protocol for removing this sample. On top of that all these technics need special staff for sampling, chain-of-custody and lab analysis which consum economical and time resources. In order to avoid the last problem some unsuccessful efforts have been made in order to build a monitor of tritium with LSC [36]. In any case, other problems still remain.

The ionization chamber (IC) is based on a gas chamber (sample) which contains electrodes connected to different voltage. This electrodes recover the ionization current that is produced due to the β radiation. It is a simple and fast system, but the problem is that on the one hand it has too high LDL, more than 10 kBq, and, on the other hand, it needs the state of the sample to be gas or steam [37] [38].

The calorimetry is based on the measurement of the heat generated due to the tritium radiation [39] [40]. The problem with this technic is that it has a too high LDL, of the order of GBq, and it needs too long time, more than 2 days, for taking a measurement.

The Beta Induced X-ray Spectrometry (BIXS) is based on the measurement of the bremsstrahlung with PMTs of NaI [41] [42] or with Silicon Drift Detector (SDD) [43] produced due to the tritium radiation. The problem with this technic is that it has too high LDL, of the order of MBq.

There are many more different methods for tritium detection, although they are less used or less experimentally developed, each one with their own problems for our objective. For example, APD [44], which we cannot use in our case because they cannot function in contact with water, the mass spectrometry [45], which needs to store the sample several months before taking the measurement or Cavity ring spectroscopy [46], which requires a special optical configuration that is not possible outside

the laboratory.

We have to keep in mind that all these techniques are offline methods that take too long to finish the process of taking measurements which include sample taking, sending the sample to the lab, analyzing of the sample so we cannot use them for tritium monitoring. LSC is the only technic which has a LDL enough low to verify the compliance with the established limit, 100 Bq/L. Therefore we will explore this area but, in order to avoid the problems related with this technic (off-line results, no-reusable liquid scintillator and the chemical toxic wastes) we will delve in solid scintillators. There are several studies that have been done so far which intend to do the same as we want with this project, to create a quasi-real time monitor of low tritium activities in water based on solid scintillation:

- First study was done by M. Muramatsu, A. Koyano and N. Tokunaga in 1967 who used a scintillator plate read out by two PMTs in coincidence [47].
- The second study was carried out by the A. A. Moghissi, H. L. Kelley, C. R. Phillips and J. E. Regnier in 1969 that used one hundred plastic fibers coated with anthracene powder and read out by two PMTs in coincidence [48].
- Third study was performed by R. V. Osborne in 1969 who used sixty scintillator plates stacked read out by two PMTs in coincidences [49].
- Fourth study was done by the A. N. Singh, M. Ratnakaran and K. G. Vohra in 1985, who used a scintillator sponge read out by electronic coincidence [50][51].
- Fifth study was carried out by K. J. Hofstetter and H. T. Wilson in 1991, who did different experiments for testing different shapes of scintillator plastic like several sizes of beads, fibers, etc. The better result which Hofstetter got for solid plastic scintillator was a efficiency of the order of 10^{-3} [52][53].

| | Efficiency, η_{det} ($cps/(kBq/L)$) | Surface F_{sci} (cm^2) | Specific efficiency $\varepsilon_{det} =$ η_{det}/F_{sci} | LDL (kBq/L) |
|------------|--|---------------------------------|---|-------------|
| Muramatsu | $3.85 \cdot 10^{-4}$ | 123 | $3.13 \cdot 10^{-6}$ | 370 |
| Moghissi | $4.5 \cdot 10^{-3}$ | > 424.1 | $< 1.06 \cdot 10^{-5}$ | 37 |
| Osborne | 0.012 | 3000 | $4 \cdot 10^{-6}$ | 37 |
| Singh | 0.041 | 3000 | $1.37 \cdot 10^{-5}$ | < 37 |
| Hofstetter | $2.22 \cdot 10^{-3}$ | ~ 100 | $< 2.22 \cdot 10^{-5}$ | 25 |

Table 1.8: Results of different scintillator detector for tritium detection [30]

The results of these experiments are summarized in the table 1.8. We can see in the first column that the intrinsic detector efficiency, η_{det} , is very different in these experiences. As we know that, in this type of detectors, one of the most important factor, which affect to the efficiency, is the active surface of the plastic scintillator, F_{sci} , and we can see in the second column that it is very different in each detector, we use the specific detector efficiency (third column), in order to compare these experiments, that's, the intrinsic detector efficiency normalized to this active surface. Now we can check that, effectively, these specific efficiencies are quite similar. On top of that we can check that the better specific efficiency was obtained for Moghissi who used scintillating fibers. This is a good point which justify our choice about using of fibers like a scintillator. Finally we can see in the last column that the LDL in all these experience are more or less similar and, they are too high for our aim so developing a detector which overcome these LDL is an essential study right now in order to monitoring the tritium levels.

1.4 Tritium project and Tritium monitor

As we have seen in the section 1.3, the current technics which exist nowadays have either higher LDL than the limit established by Council Directive, 100 Bq/L, or they are a off-line method (too slow) so those methods cannot be used for tritium monitoring in quasi-real time.

As a result of these limitations appear the *Tritium* project [54], whose title is "Design, construction and commissioning of automatic stations for quasi-real time monitoring of low radioactive levels of tritium in water".

This project has been funded by Interreg Sudoe program of the EEC in the 2016 call with the reference number SOE1/P4/EO214. The purpose of this project is the development of a tritium monitor in quasi-real time. This monitor consists of a ultra pure water system, which prepare the sample before we introduce it in our detector, the tritium detector where the tritium measure will be done, the active veto and the pasive shielding which reduce the natural background of our tritium detector and several types of electronic which control all these parts of the monitor, analyze the tritium measurement and will send an alarm if the limit of 100 Bq/L is overcome.

The tritium detector is based on measurements of low energy beta radiation from the radioactive decay of tritium. For doing this task this detector consists of scintillator fibers, that we put directly in contact with water which can contain tritium. We need to put both, scintillator fibers and tritium water, in contact due to such a low mean free path of tritium electrons (table 1.5 of the seccion 1.2). Then, the photons produced on this fibers will be read out by several photosensors. The photosensors which we have tested in this experiment are photoelectron multiplier tubes (PMT) and silicon photoelectron multiplier (SiPM) arrays.

The difficulty when we try to measure tritium is to distinguish these signals from the background. This is because tritium signals are small since tritium events has low energy (\sim keV) and this is the energy range in the spectrum where there are more background counts (the lower energy in the spectrum, the more electronic noise). We will use coincidence techniques in order to reduce the counts from the background.

It is important to check the water tightness of each prototype because if the water reaches the photosensor it will be irreparably damaged. On top of that if we use high concentrations of tritium in water for laboratory tests we can contaminate this laboratory, which could be dangerous for the healthy of the workers and it could spoil measurements of future experiments.

Finally this monitor will be installed in the Arrocampo dam, Almaraz, Spain, where the Almaraz nuclear power plant release the water which is used in their cooling system, Figure 1.4. This NPP has two nuclear reactors whose type is PWR. This dam is located near the Tago river, which is the largest river in Spain, 1007 km. This river cross from Aragon (Spain) to Lisbon (Portugal) and flows into the atlantic ocean. This river is used for an important quantity of animals, plants and even humans because the water of this river is used as drinking water by the spanish and portuguese people. Therefore the international cooperation in order to maintain the quality of this water is very important.

The *Tritium* collaboration is a international group consisting of a consortium of 6 different southwestern european institution of 3 different countries: The University of Aveiro, in Portugal, The University of Bordeaux and the CNRS (Section Aquitaine-Limousin), in France and the University of Extremadura, *Junta de Extremadura* and University of Valencia, in Spain.

FOTOO TRITIUM



(a) Arrocampo dam and Almaraz Nuclear Power Plant



(b) Tago river along Spain and Portugal

Figura 1.4 – Arrocampo dam, Almaraz NPP and Tago river

Each institution has focused in the development of a different part of all this project:

- First, the Extremadura group has developed and installed the ultra pure water system with which we get water with very low conductivity, $\sigma \approx 10 \mu S/cm$. The conductivity of the water before the cleaning process is around $1000 \mu S/cm$. This clean process is very important for two reasons. On the one hand, it is important for maintaining our detector very clean, which is a critical point. On the other hand, it's important because with this process we reduce the natural background since we remove several natural radioactive isotopes that there are in this water (except tritium) such as ^{222}Rn , ^{40}K or ^{137}Cs . This system will be explained in the section 2.3.
- Second, french group has developed the passive shielding where our detector will work inside. It is based in ultra radiopure lead with very low intrinsic activity. The objective of this passive shielding is to reduce the external natural background that affects our system, for this reason we use lead. Obviously, this shielding doesn't have to affect to the measurement of our system, for this reason we use radiopure elements with very low intrinsic activity. This shielding will be explained

in the section 2.4.1.

- Third, The Portugal and Spain people has collaborated for designing, developing and building four different prototypes of tritium detector and active vetos for removing cosmic events. These prototypes and vetos will be explained in the chapter 4 and section 2.4 respectively.
- Lastly, The Portugal and Spanish people has also developed the simulations about this system. The program which we have used in this project is GEANT 4, which is a simulation package. It consists in an extensive C++ library with which we can design the geometry of our detector, the physical processes which happen there, etc. This simulation will be explained in the chapter 5.

The tritium level which we want to measure follows the ALARA principle (As Low As Possible Achievable) and to get it there are important characteristics which our tritium detector must have:

- *Compact.* This is important because in the place where this detector will be installed the useful space that we can use is finite.
- *Thin active volume and large active area.* On the one hand, we have to take into account that, as we have seen in the table 1.5 of the section 1.2, the mean free path of the β particle of tritium decay is very low so we need to work with thin active volumes. In practice, Active thickness beyond the mean free path of the tritium will only contribute to the background. On the other hand, as we have checked in the section 1.3 the efficiency of this type of detector scales with the active area so we need to design our detector with the largest possible active area.
- *High sensitivity to tritium.* We are going to work with low tritium activities so we need to reduce as much as possible the non-detected tritium events.

- *High specificity to tritium.* We need that our detector is able to distinguish the tritium signal of the signal of other radioactive elements which can be present in the initial sample.
- *Quasi-real time response.* As we have seen it is important that our sistem can work in quasi-real time in order to detect any problem as fast as possible.
- *Rugged system.* Finally, we have to take into account that our objective will be installing an automatical system which will work during a lot of years without specialized people so we need that our monitor are rugged.

In order to get the measurement in quasi-real time we need to work *in situ*, that's, we need that our detector is able to work in the same place that we take the sample. Whit the work *in site* we achieve:

- a faster monitor because we eliminates the process of taking the sample, the chain-of-custody until this sample arrive to this laboratory and the complexity which involve these tasks.
- a better monitor since if we can work *in site*, our measurements can be more frequent hence we will can identify cahnges in the activity earlier.
- a cheaper monitor because we have not only the material costs attached to the sample collection, chain-of-custody of this sample, shipping of this sample to the laboratory, etc. but we have also eliminated the costs attached to the specialized staff who are involving in these tasks. Our detector will only need frequent calibrations each time in order to ensure its correct operation.
- a safer monitor since the personal exposure dose is reduced and the changes in activity are detected fastly. On top of that we remove the possibles mistakes which can be done by specialized staff.

1.5 Work scheme

Tesis de fibras BCF-12

This thesis is build up as follow:...

The objetive of this three-phase project was design, develop and instalation and commisioning a automatical system capable of detection tritium which we find in the water that is used by the nuclear power plants for their cooling system. The initial idea was quantifying its activity in quasi-real time before discharging it into public rivers or seas.

We are speaking all the time about getting the measurement in quasi-real time, that is, in less of 10 minuts but what about the real time? It is obviously imposible because we are mesuring the activity, I mean, we are counting tritium decays in samples with low activities (supposedly less than 100 Bq/L) so we need some time in order to get enough stadistic with which we can distinguish the tritium signal to the background in our system.

In order to get the measurement in quasi-real time we need to work *in situ*, that's, we need that our detector is able to work in the same place that we take the sample. The reason of this fact is because if we need to move the sample we lose a lot of time (1 or 2 days in some cases). Furthermore if we avoid to move the samples we get a:

- faster detector because we eliminates the process of taking the sample, the chain-of-custody until this sample arrive to this laboratory and the complexity which involve these tasks.
- better detector since if we can work *in site*, our measurements can be more frequent hence we will can identify cahnges in the activity earlier.
- cheaper detector because we have removed all the cost associated with

the sample collection, chain-of-custody of this sample, shipping of this sample to the laboratory and analysis thereof there. Not only we have eliminated the material costs attached to this tasks (material, transport, etc) but we have also eliminated the costs attached to the specialized staff who are involving in these tasks. With our detector we will only need frequent calibrations each time, which we consider suitable, in order to ensure the correct operation of this detector

- safer detector since the personal exposure dose is reduced and the changes in activity are detected fastly. On top of that we remove the possibles mistakes which can be done by specialized staff who follow each protocol of this tasks.

Dividiremos este trabajo en seis partes:

1. En primer lugar, se realizará un estudio sobre las fibras centelleadoras para determinar el protocolo de manipulación para obtener un procedimiento de preparación de un haz de fibras centelleadoras con un buen rendimiento óptico.
2. En segundo, lugar estudiaremos el procedimiento de calibración de los SiPM, fundamental para el experimento. No se necesita realizar una calibración de los PMT, paso igualmente importante al anterior, ya que este trabajo fue realizado recientemente por otro componente del grupo.
3. En tercer lugar, se describirá el primer prototipo diseñado, formado por un haz de 35 fibras centelleadoras sin clad leídas por PMT, incluyendo el protocolo del proceso de llenado con agua tritiada que tuvo que ser desarrollado para cumplir los requisitos de protección radiológica y evitar contaminación accidental, y los resultados obtenidos con el mismo.

4. En cuarto lugar, se presentarán las simulaciones realizadas con el programa de Geant4 en una configuración sencilla.
5. En quinto lugar, se expondrán aspectos a estudiar en el futuro inmediato y en etapas posteriores, durante la Tesis Doctoral.
6. Se presentarán, finalmente, los resultados, logros y conclusiones alcanzadas en el desarrollo del trabajo.

For process monitoring purposes it is needed to know how much tritium is in the processed water, which is done mostly by offline methods, also involving handling of tritiated water by the operating personnel. Since tritiated water is highly toxic, it would be advantageous to have an online and inline monitor, which eliminates the need of sample taking and also speeds up the measurement process. In this thesis the investigation of a plastic scintillator-based tritium detector is presented. The main goal was to test the concept for higher concentrations as before (up to several GBq/l concentration) and investigate possible enhancements of the scintillator geometry, besides understanding the detection process. For these purposes an experimental setup consisting of standard industrial parts was designed, constructed and calibrated for HTO. Several sample chamber geometries were tested (e.g. more scintillator plates above each other, varying sample volume, etc.), and the configurations were compared in terms of sensitivity and detected pulse height spectrum. To understand the detection process in detail, a simulation was coded and compared with the experimentally obtained results. The thesis is built up as follows: First, a brief chapter introduces tritiated water, concerning properties and processing in tritium handling facilities. The next chapter presents the scintillation counting method in detail, based on the available literature. This is followed by the discussion of the previous studies in the topic of tritiated water activity measurement, then the goals of this study are defined. Chapter 4 deals with the design considerations and the technical details of the experimental setup. The next chapter presents the optimization of the setup for measurement,

then the calibration of for tritium is presented. The chapter also includes the analysis of the pulse height spectrum of tritiated water, and results of the measurements with varying sample amount and scintillator surface, all followed by discussion. Chapter 6 first presents the technical improvements done on the setup, then the results of additional measurement series and their discussion follow. In the last chapter, the simulation of the detection process is presented, together with discussion of the results. The experimentally obtained spectrum is compared with the simulation and the implications of the differences is discussed. This chapter is followed by a summary of the thesis, together with the conclusions drawn from the results.

Chapter 2

Design principles of the Tritium monitor

2.1 Detector system overview

Dar una visión general del detector. Objetivo, partes, para que sirven y donde hablaré de ellas y algo para cerrar el apartado como el objetivo final más preciso.

Reescribir la siguiente sección: This Chapter is divided in three different sections. First of all, I speak about the theoretical foundations that govern the interaction between radiation and matter. Later, I do a theoretical explanation of the behavior of a scintillator detector and their components, plastic scintillator, specifying on plastic scintillating fibers, photosensors, specifying on PMTs and SiPMs, and electronic system. Finally, I show the main parts of the TRITIUM monitor and why each of these parts is necessary. I will delve into each part in future chapters.

2.2 Tritium detector

Due to the reasons which we have discussed in the section 1.3, the type of the detector which we have developed in order to measure the tritium that there is in water samples is a scintillator detector. It consists in a chain of three main elements:

- The scintillator, that is the material in charge of detecting the tritium event. The tritium particle or a general particle (ionizing radiation) hit this material and deposits part of their kinetic energy (or all, as in the case of the tritium event) in it through ionization and excitation. Part of this energy deposited is converted in photons, generally in the visible range.

The number of photons produced carry information about the particle detected, such as its energy, type, etc.

- The photosensor, which is the part of the detector in charge of detecting the photons produced by the scintillator that reach its sensitive element (the more scintillated photons arrive to your photosensor, the better signal you have in your detector).

The most used photosensor in nuclear physics are SiPMs and PMTs which detects some of the photons produced in the scintillator and transforms it in electrons which are multiplied with a factor of around 10^6 . This millions of electrons form a electronic pulse whose properties has information of the photons that has been detected.

- The electronic system, which is the part of the scintillator detector in charge of processsing and analyzing (first analogically and then digitally) this electrical pulse of the photosensor to give us useful information about the event detected that we can understand and interpret such as a number, for instance the activity, or some kind of spectrum like energy spectrum.

In figure 2.1 we can see the scheme of a scintillation detector where the scintillator material detects ionizing radiation and produces photons that will be guided by the reflector and the light guide to the photosensor. There, some of the photons that reach the sensible part of the photosensors will be converted and multiplied into millions of electrons that will form an electronic pulse. The output signal of the photosensor (electronic pulse) will be processed and analyzed by the corresponding electronics:

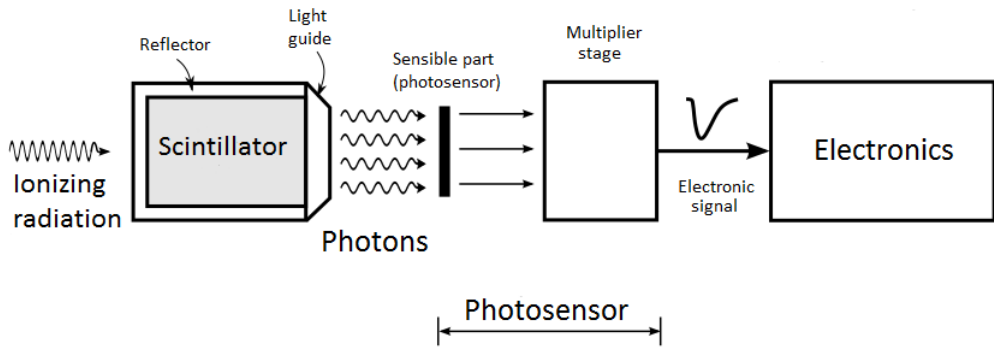


Figura 2.1 – Scheme of the scintillator detector [?]

2.2.1 Interaction of fast electrons and photons with matter

In this section, the explanation will only focus on the particles and energy range that are interesting for this thesis, which are electrons (0 – 18 keV) and photons in the visible range (approx. 380 – 750 nm).

On the one hand, electrons have charge so their interaction with matter are mainly produced with the orbital electrons that there are in that matter due to the Coulomb force. The trajectory which electrons follow is much more tortuous than other heavier particles because the mass of both interacting particles is equal, electrons. Furthermore, for the same reason, these electrons lost a significant amount of energy in each collision.

In order to speak about the total energy lost of particles in matter the specific energy loss is defined as $S = -\frac{dE}{dx}$ which expresses the energy loss suffered by the particle per unit of trajectory. In the case of electrons, this total energy loss has two main contributions, the collisions (elastic and inelastic) and radiative processes (bremsstrahlung):

$$\frac{dE}{dx} \approx \left(\frac{dE}{dx} \right)_c + \left(\frac{dE}{dx} \right)_{br} \quad [55][56] \quad \frac{\left(\frac{dE}{dx} \right)_{br}}{\left(\frac{dE}{dx} \right)_c} \approx \frac{EZ}{700} \quad [55] \quad (2.1)$$

Where E is the energy of the electron in MeV and Z is the atomic number of the absorbing material. Due to this energy loss, the electrons can only penetrate a material as far as they go before losing their total energy. This distance is known as range and, in the case of tritium electrons, its value is seen in the table 1.5.

On the other hand, photons don't have charge. Its possible interactions with the matter are photoelectric effect, Compton effect, coherent

scattering and pair production and the probability of each process depends on the energy of the photon, $E_\gamma = h\nu$, and the atomic number of the material, Z , as you can see in the figure 2.2.

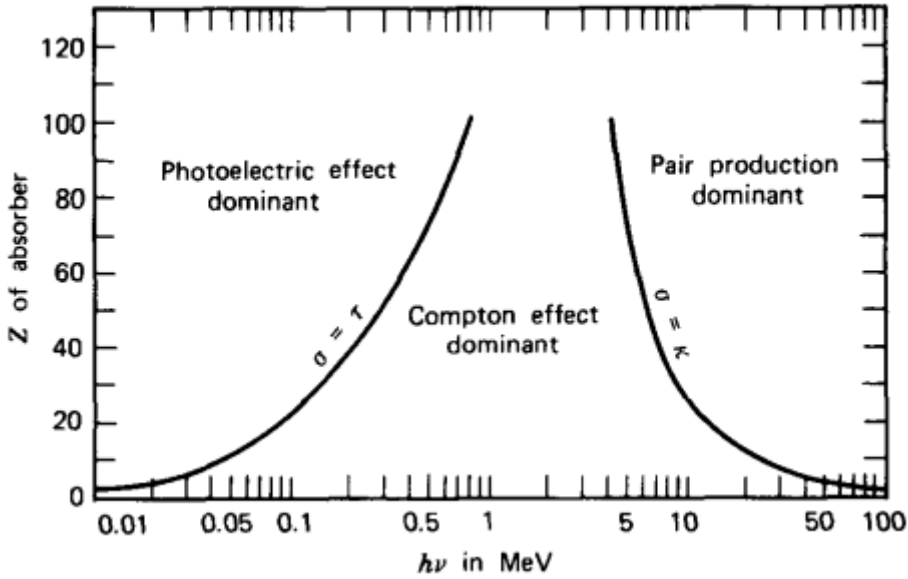


Figure 2.2 – Domain regions of the three most probable types of interactions of gamma rays with matter. The lines show the values of Z and $h\nu$ where the two neighboring effects are equally likely. [55] [56]

We have to take into account that the only relevant photons for this thesis are in the visible range, between 400 and 700 nm, that corresponds with energies of the order of the eV. Therefore the last effect, pair productions, will be not explained here because it needs a photon energy equal or more than 1.022 MeV for happening and it is not our case.

The photoelectric effect occurs when a photon interacts with an orbital electron in the material, losing all its energy. This energy is absorbed by the electron that is released from the atom (ionization). The energy of the resulting electron, E_e , is:

$$E_e = E_\gamma - E_b \text{ [55][56]} \quad (2.2)$$

Where E_b is the binding energy of the electron in this material. The probability of this effect depends on the number of available electrons through the variable Z , and the energy of the electron according to the following expression:

$$(Pr)_{Ph-eff} \approx \frac{Z^n}{E_\gamma^{3.5}} \text{ [55]} \quad (2.3)$$

As we can see in this expression and in the figure 2.2, the photoelectric effect is most probably if we use elements with high atomic number. This is the reason why elements with high atomic number are the best insulators against gamma radiation and this is one of the reasons why we use lead ($Z = 82$) for building our passive shielding as we will see in the section 2.4.1. This is also the reason why elements with high atomic number are used in the cathode of PMTs.

The Compton effect occurs when the photon interacts with an orbital electron of the material, transferring part of this energy to the electron, which is released, and this photon is scattered at an angle θ with respect to the original direction. If we neglect the binding energy, the energy transferred to this electron, E_e , is shown in the following equation:

$$E_e = \frac{\frac{E_\gamma^2}{m_0 c^2} (1 - \cos\theta)}{1 + \frac{E_\gamma^2}{m_0 c^2} (1 - \cos\theta)} \text{ [55][56]} \quad (2.4)$$

Where m_0 is the rest mass of the electron and c is the speed of the light in the vacuum. The probability of the Compton effect is proportional to the atomic number (more available electrons), Z , and decreases with the

energy of the photon.

As we can see in the figure 2.2, in the energies of the photons belonging to the visible range of the electromagnetic spectrum (of the order of eV), the Compton effect is only more likely in very light materials, ($Z < 4$). For heavier materials the photoelectric effect is the dominant effect.

Finally, in the coherent scattering, the atom is neither excitation nor ionization and the photon conserve all their energy in this collision. This effect is more probably for photons with low energies and materials with high atomic numbers. Because of the fact that the energy of the photon doesn't change we will not speak more about this effect but it is important since this effect change de direction of photons and it will affect to their mean free path.

2.2.2 Scintillators plastic

The use of scintillators in radiation detection is one of the most used technics in nuclear physics. The scintillator is a material that is able to convert the kinetic energy of the incoming particles in light (photons in the visible energy range) which we can detect and quantify. It happens because the radiation excites and ionizes the scintillating atoms which, after that, are immediately de-exciting (with times of the order of picoseconds), emitting photons.

This conversion should be linear in a wide energy range of incoming particles and it is necessary that this material has good optical properties, such as being transparent to the wavelenght of their own emission and having a refractive index as close as possible to the glass for optimizing optical coupling with photosensors.

The photon emission in the scintillator is a stadistical process,

which means that two exactly the same events will emit different number of photons. It follows Poisson statistics so when we speak about the number of emitted photons we speak about the mean number of photons.

There are two types of scintillators, organics and inorganics. Inorganic scintillators normally have a higher atomic number and density so their light output is higher. Due to these reasons they are better for gamma-ray spectroscopy (take into account figure 2.2). Organic scintillators are generally faster and they are commonly used for beta spectroscopy and neutron detection. In this section I focus my explanation mainly on organic scintillators since they are the ones we have used in our research.

Organic scintillators are based on a scintillator material, which produces light, dissolved in a base solvent. This solvent is normally based on aromatic hydrocarbons (carbon atoms linked together), that is, they are mainly composed of carbon and hydrogen atoms as we can see in the molecules of some of the most widely used scintillators, $C_{18}H_{14}$, $C_{24}H_{22}N_2O$ or $C_{15}H_{11}NO$ whose average atomic numbers are between 3.5 and 5.

The scintillator molecules, in which the organic scintillators are based, have the so called π -electron structure. The energy levels of their electrons are commonly illustrated with a Jablonsky diagram, figure 2.3, where we can see the fundamental single state, S_{0i} , where the valence electrons are, the excited single states, S_{jk} , and the excited triple states, T_{lm} . The energy difference between S_1 and S_0 states is around 3 or 4 eV, energy range of the visible photons. We can see in this diagram that each of these energy states are subdivided in smaller energetic sublevels whose distance between them are around 0.15 eV. This finer energy structure is due to the excitations of molecular vibrational modes and they are expressed with the second index of the energy states.

Because of the reason that the distance between all energy levels and sublevels are larger than the thermal energy, 0.025 eV, non-exciting elec-

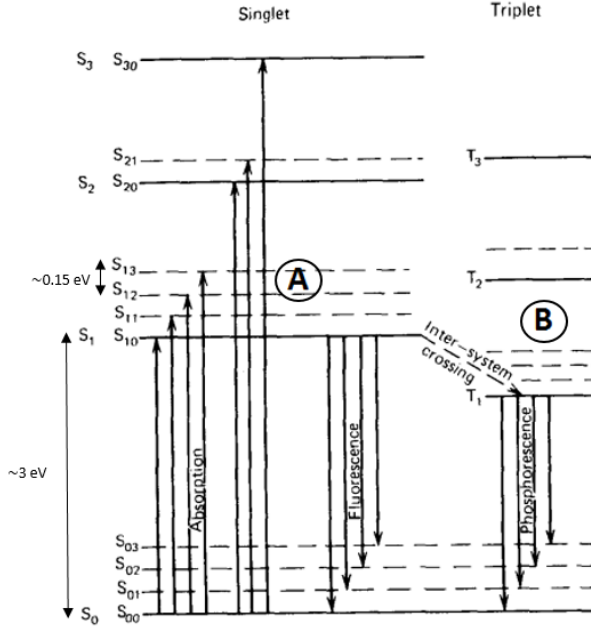


Figure 2.3 – Jablonsky diagram. [55]

trons are in the lowest state S_{00} at STP (standar temperature conditions).

When a particle deposits their kinetic energy on a scintillator, their valence electrons are exited to higher single energetic states very fast (times of the order of picoseconds) which is expressed with upwards direction arrows in the figure 2.3 and they are quickly de-excited to the first single excited state, S_{10} , through non-radiative processes known as internal conversion.

Now, this electrons can de-excited to the fundamental single state, S_{00} , through three different physical mechanisms:

- The prompt fluorescence(process A in figure 2.3), where the electron in the S_{10} energy level is de-excited to some sublevel of the fundamental state S_{0i} , emitting a photon with an energy equal to the energy

difference of these levels (around 3 or 4 eV , visible light). This process happens immediately after the excitation of the scintillator molecules (around tens of nanoseconds after excitation). Each scintillator has a characteristic emission spectrum that defines its response due to the fluorescence mechanism.

Now we can understand why organic scintillators are practically transparent to their own fluorescence emission. This is because of the reason that there exist a quenching effect in each de-excitation process whereby there are a lost of radiated energy. Due to that, all emitted photons by the scintillator have less energy than the required energy for excitation. This effect is called Stokes shift and it's represented with a general wavelength spectrum in the figure 2.4.

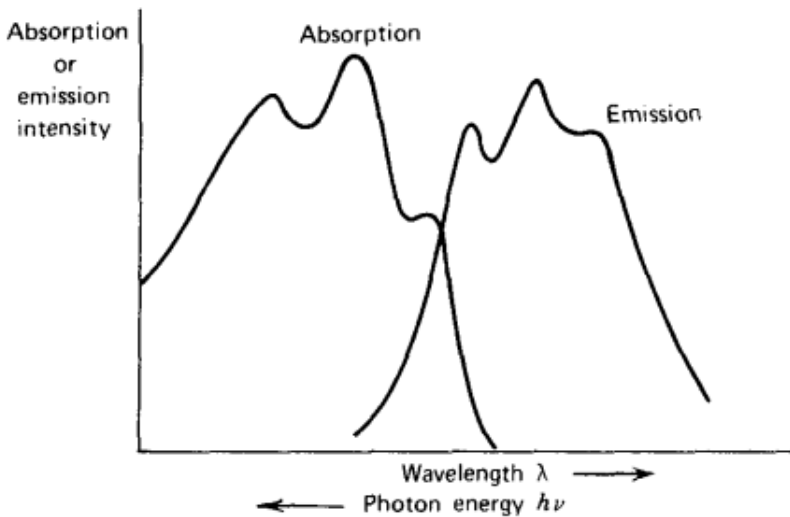


Figure 2.4 – Stokes shift. [55]

One of the most important parameters in nuclear physics is the scintillation yield, which is the fraction of particle energy that is converted in light. All mechanisms which don't produce prompt fluorescence, like phosphorescence or delay fluorescence, which we will see later, or even internal conversion, contribute to reduce this parameter. The

signal of the scintillator depends on the type of particle so, the scintillation yield will be different. This factor is normally given by the manufacturer for mips (minimum ionized particles, that's, for example, electrons with 500 keV or more) in number of photons per MeV. The intensity of the fluorescence emission in an organic scintillator during time is a combination of two exponential functions, one associated with the lifetime of the level, τ (on the order of nanoseconds), and the other associated with the energetic level population, τ_1 (on the order of picoseconds).

$$I = I_0 (e^{t/\tau} - e^{t/\tau_1}) [55] \quad (2.5)$$

- The phosphorescence, where the electron that is in the first single excited state cross to a triple excited state (process B in figure 2.3) with a process called "intersystem crossing". This is a metastable state with a much longer lifetime so electrons in this state are de-excited to the S_{0i} state, emitting a photon much later than phosphorescence. This process can happen up to 10^{-3} seconds after scintillator excitation.
- Delayed fluorescence, which occurs when an electron is in a triple excited state but whose transition to the ground state is forbidden. In this case, this electron can interact with another that is in a similar state and return to the first single state (S_1 is a more energetic level than T_1) and quickly de-excited to the ground state.



This emission has the same emission spectrum as immediate fluorescence, but occurs later.

The scintillating detectors generally use the prompt fluorescence light as output signal so a good scintillator should increase it and reduce other possible physical mechanisms.

Plastic scintillators are organic scintillators that has been dissolved in a solven and polimerized. They are easy to machine and can take any desired shape during construction. Among the forms most used today we can find blocks, thin sheets, cylinders, etc.

In our experiment we have been working with an plastic scintillator in the form of fiber, specifically, commercial fibers BCF-12 from Saint-Gobain Crystals Inc [57]. This type of fiber was chosen as the result of a comparative study among some of the best-known commercial manufacturers, such as Kuraray.

The BCF-12 fibers consist of a scintillated core, whose material is polystyrene, one of the most used solvents for plastic scintillators [55], with the possibility of surrounding it of a cladding of polymethylmethacrylate (PMMA) (smaller refractive index than core in order to archieve a critical angle) or a multicladding (second cladding) with even smaller refractive index.

When a particle deposits all or part of its kinetic energy, some photons are produced in the fiber core as a result of the scintillating process. The quantity of photons which are produced depend on the scintillation yield, whose value is around 8000 photons per MeV for a mip in our case (BCF-12 fibers) . It means that, for instance, for tritium electron, this fibers will release a maximum of around 148 photons (when tritium electron has the maximum energy, 18.6 keV), probably less because electrons with these energies are not mips.

These photons will shape the useful part of the response of the scintillator (fluorescence) for us. The energy (or wavelength) of these scintillated photons follows the distribution of their emission spectrum which, for the used fibers, is shown in the figure 2.5.

After the production of scintillated photons, we need to guide these photons to the sensitive part of the photosensor where we will detect them

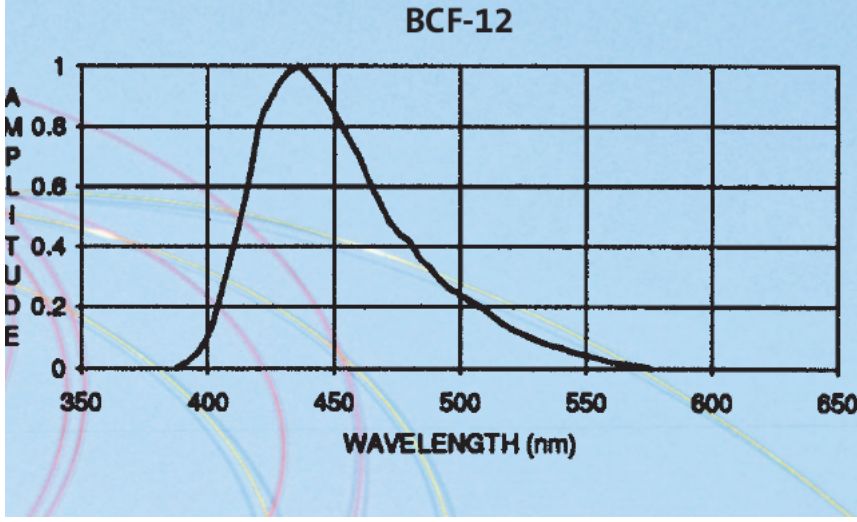


Figure 2.5 – Emission spectrum of BCF-12 fibers of Saint-Gobain. [57]

with some probability. Fibers (and scintillators in general) use the optical property of Snell's law [58] to guide their photons to the desired part (the ends of the fibers). It is based on the interface created between the core and the surrounding material. When a photon hits this interface, it is refracted (and therefore lost) following the Snell equation, 2.7. If the surrounding material has a lower index of refraction than the core of the fiber, there exist a critical angle, θ_c , at which, for angles equal or larger than this one, the photons will be totally refracted (and therefore conserved in the fiber for being guided). This effect is showed in the figure 2.6.

$$n_0 \sin(\theta_0) = n_1 \sin(\theta_1) \longrightarrow \theta_c = \arcsin\left(\frac{n_1}{n_0}\right) \quad (2.7)$$

There exist a parameter which define the efficiency of the scintillator in order to guide photons, the trapping efficiency. For BCF-12 fibers is between 3.44% and 7% (depending if the event was detected near the fiber axis (minimum) or near the core-clad interface(maximum)).

Therefore, from these 148 photons initially created with the tritium electron with the maximum energy, only a maximum of around 10 photons (for maximum trapping efficiency) will arrive to our photosensor. As you can see, we work with very weak detector signals, where there is more electronic noise and, as you will see in future chapters, we have made a big effort to reduce this electronic noise as much as possible with several technics.

In the figure 2.6 we can see how a scintillating fiber works.

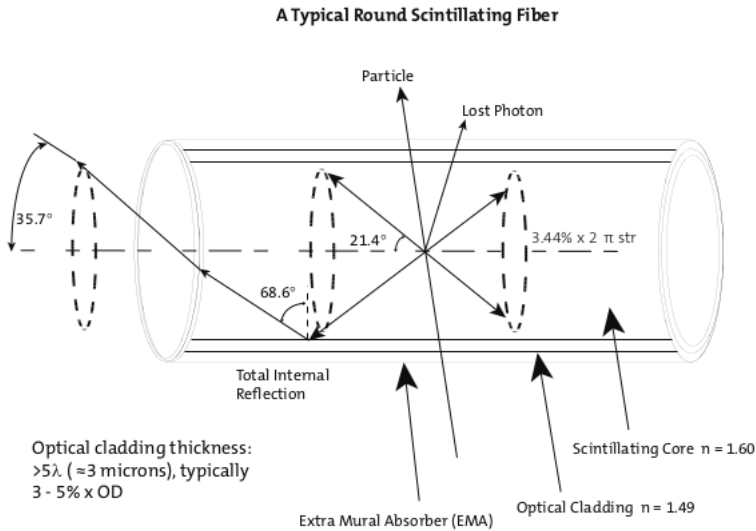


Figure 2.6 – How photons are collected in a fiber with single clad. [57]

The cladding material is useful for protecting the core surface from dirt or aggressive external agents that can reduce the light collection but at the cost of losing some light because it increase the critical angle. In the table 2.1 we have three different examples where this effect is illustrated.

This is what theoretically happens but, in the practice, it's difficult to archieve a perfect air-core or water-core interface and it will affect to the light collection. Due to the reason that the commercial claddings are thicker ($30 \mu\text{m}$) than the mean free path of tritium in water (around $5 \mu\text{m}$) we cannot use commercial cladding in our detector hence we will need to take

| Material | Refractive index | critical angle ($^{\circ}$) |
|------------------|------------------|-------------------------------|
| Air | 1 | 42.98 |
| Water | 1.33 | 62.47 |
| Cladding of PMMA | 1.49 | 76.26 |

Table 2.1: Critical angles asociated to different interfaces created with polystyrene, $n_0 = 1.6$, and other materials

special attencion for archieving a water-core interface enough good. As we will see in the section ??, we have used a special protocol developed in the ICMOL (pie de pagina explicando que es el ICMOL) for preparing fibers before we use them for tritium detection.

Some of the most important properties of the used fibers are summarized in the table 2.2.

| | |
|--|---|
| Core material | Polystyrene |
| Core refractive index | 1.60 |
| Density | 1.05 |
| Cladding material | Acrylic (PMMA) |
| Cladding refractive index | 1.49 |
| Cladding thickness (μm) | 30 |
| Numerical aperture | 0.58 |
| Trapping efficiency | 3.44% minimum |
| Radiation lenght (cm) | 42 |
| Emission peak (nm) | 435 (Blue) |
| Decay Time, (ns) | 3.2 |
| 1/e Length (m) | 2.7 |
| Scintillator yield ($\#\gamma/\text{MeV}$) | ~ 8000 |
| Operating Temperature | -20°C to 50°C |

Table 2.2: Properties of BCF-12 fibers from Saint-Gobain Inc. [57]

2.2.3 The light detection in photosensors

So far we have created the scintillating photons in the core of the fiber, which have been guided to its ends. Now, what we need is the so-called photosensor, which is an element that is able to detect these scintillating photons. Photosensors have a sensitive part that is optimized to detect photons in a range of energy (normally inside of a visible range) with enough efficiency. After that, the photosensors create an electronic signal that carries information about these photons detected, such as their number or their detection time.

One of the most important things in the scintillation detector is that the emission spectrum of the scintillation (figure 2.5 in our case) overlaps as much as possible with the detection efficiency spectrum of the photosensor used, specifically their higher peaks. In this case, the efficiency of this detector, which is proportional to the multiplication of both factors at the same photon energy, will be optimized (the largest).

There are a lot of different photosensors that can be used for this purpose, whose photon detection relies on totally different physical processes, such as photoelectron multiplier tubes (PMTs), silicon photoelectron multiplier (SiPM) or charge-coupled device (CCD). Each one of these will have different properties and we have to choose the one which fit better for our objective.

Our main proposal for our scintillation detector will be to use SiPM arrays because they are very fast (of the order of ns) and have high photodetection efficiency (a maximum of around 50%) and high gains (multiplication factor of 10^6) with a low voltage supply. On top of that, one of the most important reason of this choice is that SiPM arrays are able to detect a single photon with high efficiency, which is very important since, as we have seen in the section 2.2.2, just a few photons will arrive to the sensible part of the photosensor. We will test also the PMTs, which are the

conventional choice, because they are still interesting since they have lower dark count rate than an equivalent SiPM and some similar properties like its gain.

Photoelectron multiplier Tubes (PMTs)

Photoelectron multiplier tube is one of the most used photosensors in nuclear physics during last decades. Its main objective, like all photosensors, is to detect the scintillating photons that reach its sensible part and convert it in an electronic signal large enough to be measured.

In the figure 2.7 we have a schematic drawing where we can see the PMT components and how it works. First of all, as we can see in the figure 2.7, we need to create electrons that will travel in the medium (electronic signal). To increase the amount of conserved electrons, we need to work inside a vacuum tube. Therefore, the PMT consists of a vacuum tube that has a glass window through which photons will penetrate inside.

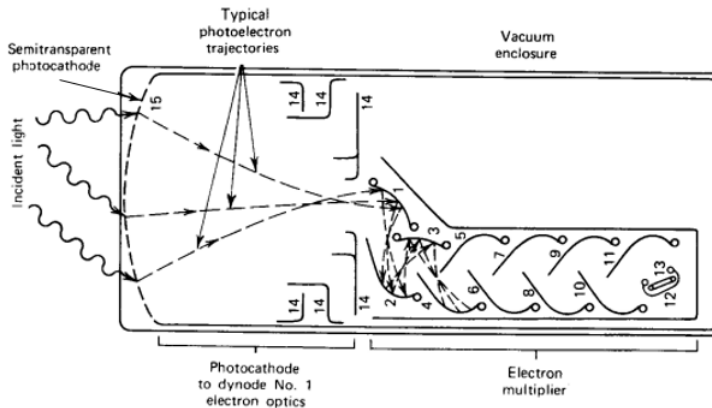


Figure 2.7 – Scheme of a PMT. [55]

The way in which PMT achieves their aim of detecting scintillating photons happen inside this vacuum tube and it is based in two different phases:

- First, the PMT convert photons that reach its sensible part in electrons, called photoelectrons, with some probability through photoelectric effect. This sensitive part is the photocathode, which consists of a thin layer (thickness of the order of nanometers) deposited on the inner surface of the PMT windows. The material of the photocathode is chosen to increase the probability of producing photoelectric effect with the scintillating photons. The PMTs which we have used in this experiment are the model R8520-406 from Hamamatsu [60]. The material of the photocathode in our case is Bialkali.

The response of the PMT at long wavelengths is limited mainly because the photon does not have enough energy to produce a photoelectric effect or the emitted photoelectron does not have enough energy to overcome the material-vacuum interface. The response of the PMT at short wavelengths is limited mainly due to absorption in the window material. Due to both reasons, the response of the PMT will have a strong dependence with the energy of the photon and it's commonly expressed in the quantum efficiency (QE) spectrum which is the quotient between the number of photoelectrons produced at the cathode of the PMT and the number of photons reaching it. For our PMTs, is showed in the figure 2.8.

The maximum values of the PMT quantum efficiency is commonly between 20% and 30% [55] (a little bit less than 30% in our case [60]). If we compare the emission spectrum of our scintillating fibers, figure 2.5, with the quantum efficiency spectrum of the PMTs that we have uses, figure 2.8, we can see that the peaks of the spectrum are more or less in the same position (435 nm for the fibers and 420 nm for the PMT and approximately the same value for 435 nm). As I said in the subsection 2.2.3, it is very important for increasing the overall efficiency of our scintillation detector.

- Next, Due to the reason that the number of photoelectrons in the photocatode is very small, we need a electron multiplication stage

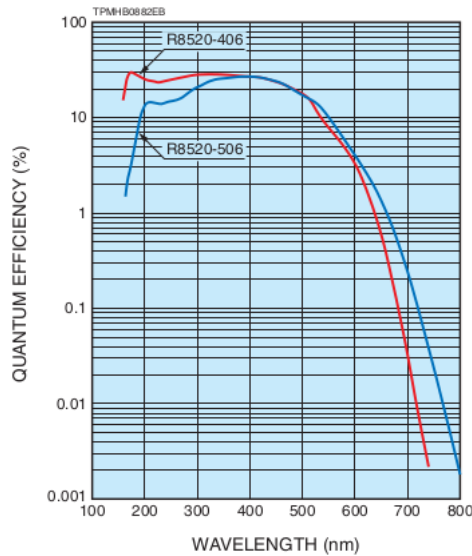


Figure 2.8 – Quantum efficiency spectrum for the PMT used (R8520-ZB277). [60]

to achieve a large enough electronic signal to be processed by the electronic system.

This stage is based on three elements, focusing electrodes, dynodes and anode: They are metallic sheet whose shape and position are designed to optimize the collection of the electrons. The PMT needs a high voltage (HV) which are distributed between all this elements, including the photocathode, in a increasing voltage way in order to attract and accelerate the electrons. This voltage division is achieved with an electronic circuit than can be fed with positive voltage (ground in the photocathode) or negative voltage (ground in the anode). The comercial electronic circuits of Hammatsu are showed in the figure 2.9.

The electronic circuit that can be supplied with negative voltage is faster due to the ausence of the capacitances C4 and C5, but the other circuit, supplied with positive voltage, can be interesting for other tasks such as measurement of PMT currents. We will use both,

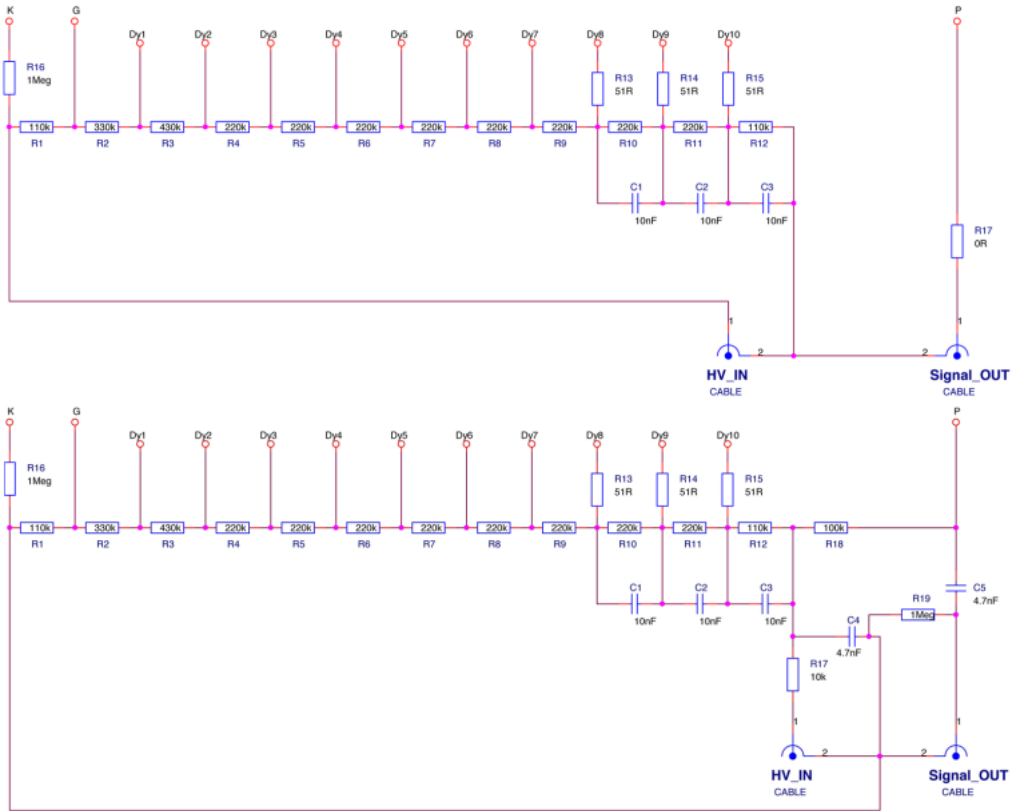


Figure 2.9 – Hamamatsu commercial voltage divider electronic circuit. Upper circuit with negative supply and lower circuit with positive supply. [60]

depends on the objective of the study.

Focusing electrodes are used to get the photoelectrons to reach the first dinode. Therefore, they have an collection efficiency (CE) that is defined as the quotient between the number of photoelectrons reaching the first dinode and the number of photoelectrons leaving the photocathode and whose value is around 80% for PMTs.

The dynodes is the part where the multiplication takes place. They have different voltage between each dynode in order to accelerate the electrons sufficiently so that, when the electron hit the each dynode, several electrons are emitted. The multiplication factor, δ , is the multiplication of electrons in each dinode and its value is commonly around 5 and it has a strongly dependence with the HV. If we take into account that the PMT has N dynodes, commonly N=10 and we guess that each dynode has the same gain, δ , the overall gain of the PMT is:

$$G = CE \cdot \delta^N \quad [55] \tag{2.8}$$

If we use the numerical values mentioned in this section, we can see that the overall gain of a general PMT will be of the order of 10^6 . It is important to mention that this value depend strongly on the HV used.

It is important to note that the uncertainty of the output signal with this gain is much greater than without it. That is why there are times when it is interesting to work without gain, such as when we try to count the number of photons. We can achieve it with a small modification of the electronic voltage divider circuit 2.9. It consists of short-circuiting all the dindes and the anode. In this way we are collecting the signal in dinode 1 in which no amplification has occurred. We will use this voltage divider circuit without gain in our study with fibers.

Finally, the anode is the point where the collection of all the electrons produced in the multiplication process takes place and it is the one that gives rise to the output signal of PMTs.

Due to the fact that all intermediate factors (photoelectric effect and multiplication of electrons) are linear, the output signal of a PMT will be linear with the number of photons that reach its sensitive part. This output signal has a spread of the order of tens of nanoseconds.

The multiplication of electrons can be described as a Poisson statistical process, so, for each electron in the first dinode, we will have G new electrons whose variance will be approximately \sqrt{G} .

Finally, we have to take into account that the photocathode can emit electrons whose origin doesn't belong to the scintillation light. This signal, which is named dark current, can happen due to several reasons like cosmic radiation, light from environment or thermoionic emission (the dominant) and this value will be around 2 nA according to Hammamatsu data sheet.

The calibration of the most important parameters (for our objective) of the PMTs used, which are dark current, gain for several HV and quantum efficiency, have been done at IFIC in the framework of NEXT experiment [59].

Silicon Photoelectron Multiplier Array (SiPMs array)

Basado en la tesis de NEXT y completar con cosas del TFG de Andrea Chana y la tesis larga.

Me quedo con el SiPM array de area de SiPM mayor debido a la sensibilidad... punto 4.2 de la tesis de SiPM (LARGA).

resumen de como varía cada magnitud con la temperatura y el voltaje. Tesis SiPMs.

tabla con las propiedades del SiPM.

Hablar de la electrónica como en el tema 5 de la tesis larga.

Incoherent light source!

fired cells.

Trabajo de Fernando Hueso.

Con la serie S13360 de SiPMs (la nuestra) -> The MPPCs inherits the superb low afterpulse characteristics of previous products and further provide lower crosstalk and lower dark count.

Describir un APD -> Tesis NEXT. Describir un SiPM -> Tesis NEXT, completar con lo de Andrea y la tesis larga. Descripción de SiPMs array. Data sheet

SiPMs are small solid state photosensors with signal levels similar to PMTs and excellent photon counting capabilities. Their use as imaging cells in NEXT-100 has moreover the advantage of providing an optimal spatial resolution (active area down to 1 mm^2) at a moderate cost and a reduced radioactivity budget (micro Bq/kg level for the ^{238}U and ^{232}Th radioactive chains [21]). SiPMs have an optimal sensitivity in the blue spectral region (around 440nm).

B. Photon Detection Efficiency -> entero.

Algunas de estas ventajas son: inmunidad a campos magnéticos, facilidad de uso (simpleza de resultados generados), tamaño compacto, buena linealidad, electrónica simple, alta ganancia a bajo voltaje de polarización, alta eficiencia de detección de fotones (Photon detection efficiency),

tiempo de respuesta corto, alto ritmo de conteo (count rate), buena resolución temporal, y un amplio rango de respuesta espectral (MANUAL MPPC HAMAMATSU, España et al., 2009, Ramili et al, 2012). Se ha comprobado que estos aparatos tienen buen desempeño bajo campos magnéticos de 0 a 7 Tesla, al igual que su conveniencia para adquisiciones tipo PET bajo la presencia de gradientes cambiantes y secuencias de radiofrecuencia de MRI (España et al., 2009, p. 1)

Aunque el aumentar el voltaje inverso mejora la eficiencia de detección de fotones, también aumenta la corriente oscura. Reducir la temperatura sin embargo disminuye la corriente oscura.

Para contar el número de veces que dos o más fotones son detectados simultáneamente, el umbral es establecido en $N - 0.5$ p.e. (donde N es un número arbitrario de fotones). Al contar el número de pulsos que exceden este umbral se puede saber el número de veces que se han detectado simultáneamente N o más fotones (Manual Hamamatsu, 2008, 2007)

En este trabajo se estudió tanto la variación en la resolución en energía de un SiPM como la variación del centroide de un pico (explicado a detalle en los próximos capítulos) con la temperatura y el voltaje. Para ello se estableció una electrónica de adquisición adecuada para la mayor eliminación de ruido posible y óptima resolución (explicada en el Capítulo 3).

Superponer 2 plots con el LED a distintos voltajes (2 o mas... probar varios voltajes a V_{ov} recomendado y quedarnos con los mejores).

Intro 1.1 de la tesis "Uso de fotomultiplicadores de silicio para medidas de alta velocidad y baja intensidad luminosa".

"Solid state technologies have evolved a lot in recent decades and the development of new high sensitivity semiconductor devices is a reality nowadays. In fact, current Silicon Photomultipliers (SiPMs) are replacing

PMTs in many fields, and probably, they will not have rival in a next future."

Basarse en la memoria de Andrea Chana y completar

Un fotomultiplicador es un aparato que convierte señales luminosas en señales eléctricas.

poner PDE en la lista de nomenclatura

poner APD en nomenclatura

Durante los últimos años se han introducido fotodetectores de estado sólido compatibles con altos campos magnéticos: APDs y más recientemente SiPMs. Esta tendencia de uso de detectores compactos, unido al aumento del número de canales en los detectores, ha hecho imprescindible el desarrollo de electrónica integrada en PET. La arquitectura electrónica típica para PET se basa en la digitalización y lectura de la carga que proporcionan los fotosensores, para lo cual se realiza un proceso de amplificación de bajo ruido y adaptación de señal al digitalizador. Sin embargo, existen hoy en día otras formas de detección alternativas, como esquemas time over threshold (ToT) o digitalizadores directos de formas de onda.

Los detectores basados en semiconductores, que comprenden a los APDs y a los SiPMs, son insensibles a los campos magnéticos y actualmente ofrecen una alternativa a la opción clásica basada en PMTs. Ambos dispositivos generan portadores libres debido al efecto fotoeléctrico, produciendo una corriente eléctrica proporcional a la intensidad de la luz incidente. Cuando los fotodetectores son capaces de multiplicar fotoelectrones primarios, se pueden producir corrientes eléctricas grandes incluso en el caso de que incida un único fotón.

Figura 11 de la tesis de cristales monolíticos

Este proceso multiplicativo en un APD también produce un ruido estocástico inherente que no está presente en un fotodiodo convencional y

que se conoce con el nombre de factor de ruido en exceso (F). Este ruido aumenta según lo hace la ganancia M del APD (Ec.15), lo que obliga a operar con una ganancia interna moderada, ya que este ruido en exceso es uno de los principales factores limitantes con estos dispositivos.

La ganancia M , depende exponencialmente de la tensión de polarización inversa del dispositivo (Fig. 11, derecha). Sin embargo, en la región de operación de los APDs con $M \approx 100$, un cambio relativo de tensión de polarización corresponde a un cambio lineal en la ganancia, con pendientes típicas de 10 %/V. Además, la temperatura debe estar debidamente estabilizada en un sistema con APDs ya que estos detectores sufren variaciones importantes con la temperatura, típicamente $\approx 2-3\%$ / °C

Todo el apartado de SiPMs de esta tesis. Lo de antes era de APDs.

IMPORTANTE PARA JUSTIFICAR EL SIPM ARRAY ESCOGIDO
-> 4X4:

El número de celdas de un SiPM dependerá de la aplicación específica. Será lo suficientemente elevado para detectar la cantidad de fotones esperada pero sin exceder innecesariamente este valor, ya que cada celda necesita de espacio para las resistencias de quenching de cada APD y para la separación y aislamiento entre las diferentes celdas. Cuanto mayor sea el número de celdas, mayor será el espacio muerto y menor su eficiencia. Por el contrario, un número de celdas inferior con celdas de mayor tamaño, implica una alta eficiencia de detección de fotones (Photon Detection efficiency, PDE) pero un rango dinámico bajo. La PDE en un SiPM se define como su eficiencia cuántica por el ratio entre el área sensitiva y el área total del dispositivo, lo que se conoce como factor de llenado (fill factor) y se representa con “epsilon”, por la probabilidad de que un fotoelectrón comience un proceso de avalancha (Ec. 19). Aparte de la longitud de onda,

señal lineal con la luz entrante.

MPPC Multi-Pixel Photon Counter

Linear response with the incoming photons.

Ver apuntes generales comentados de fotosensores en la intro de fotosensores

Los tubos fotodetectores (PMT) son los dispositivos adecuados para esto, aunque, sin embargo, los avances tecnologicos de las ultimas decadas en tecnologia de semiconductores ha permitido el desarrollo de los fotodiodos de avalancha operados en modo Geiger (APD, Avalanche PhotoDiode)

Actualmente, la tecnologia de semiconductores ha permitido el desarrollo de fotodiodos que, por sus prestaciones, pueden reemplazar adecuadamente a los PMT.

4.5. Recent trends on SiPMs The size of commercially available SiPMs is currently limited to approximately 10 mm² [164] Roncali-2011 . The restricted are is dictated both by manufacturing and noise considerations.

Most of the devices produced up today have an area of 1 x 1 mm² . SiPMs of 3 x 3 mm² and 4 x 4 mm² have been also produced, but their use is limited since the noise level scales with the area of the device. 2D matrixes of 1 x 1 mm² SiPMs devices on the same substrate are suitable. Such devices increase the detection area maintaining on each read-out channel a noise level corresponding to a SiPM of 1 mm² and providing at the same time 2D position information with a spatial resolution of 1 mm² [165] Dinu-2009 .

produced in the process. If each ionization process could be considered independent of the others, the fluctuations would then be described by a Poisson distribution where the variance (σ^2) would be equal to

the mean number of ionization electrons, N_I . However, the fluctuations in the mean number of ionization electrons present a lower value, as predicted by Fano's theory [66], being proportional to a factor F , known as the Fano Factor, which multiplies the mean primary ionization yield.

NEXT: To shield NEXT-100 from the remanent external flux of high-energy gamma rays, a lead castle structure has been designed and built (see Figure 3.9). The lead wall has a thickness of 20 cm and is made of layers of staggered lead bricks held with a steel structure. The lead bricks have standard dimensions (200 x 100 x 50 mm³), and by requirement, an activity in uranium and thorium lower than 0.4 mBq/kg. The lead castle, with a total weight of 60 tons,

Por su parte, el fotomultiplicador de Silicio (SiPM en adelante), desarrollado a finales del pasado milenio, acomete finalmente con éxito el reto de detectar, medir temporalmente y cuantificar señales poco luminosas hasta el nivel de un solo fotón. Los SiPM ofrecen una alternativa altamente atractiva que replica minuciosamente las propiedades de detección en condiciones de baja luminosidad de los PMT, proporcionando al mismo tiempo todos y cada uno de los beneficios de los detectores de estado sólido. Un SiPM ofrece la opción de trabajar con bajos voltajes de operación (menos de 100 V normalmente), posee insensibilidad a los campos magnéticos, destaca por su robustez mecánica y excelente uniformidad de la respuesta y es además muy compacto en dimensiones.

Comparison of photosensors considered

2.2.4 Readout system of photosensors

Ver tesis de detectores monolíticos.

Electronical system for PMTs

2 posibles configuración. Cuando leo con un solo PMT... Cuando leo con 2 PMTs en coincidencia...

Esto es así para todo, vetos, Tritium prototypes, etc...

Calibración de PMTs...

Electronical system for SiPMs arrays (PETSYS)

Ver pagina 82 de la tesis de detectores monolíticos

Ver hoja resumen entre los distintos tipos de ASICS comerciales, página 82 de la tesis de detectores monolíticos. Plantearlo como que elegimos PETSYS porque es mejor respecto a los otros.

Leer el tema 3 (parte de electrónica) y tema 4 (parte del flexToT)

2.3 Ultrapure water system

2.3.1 Introduction to the ultra pure water system

2.3.2 Set up of ultra pure water system

2.4 Tritium monitor background shields

2.4.1 Set up of the passive shield (lead)

2.4.2 Set up of the active shield (cosmic veto)

Chapter 3

Research & Development on detector design and components

3.1 Introduction (why is important)

Porque estamos haciendo este estudio? CUales son las grandes preguntas que pretendemos resolver en este capítulo (no en el experimento si no en este capítulo). Porque lo estamos haciendo? Como lo hemos hecho? En las conclusiones decir si se ha conseguido o no

3.2 Characterization and R&D on scintillating fibers

3.3 Characterization and R&D on SiPM arrays

3.4 Characterization and R&D on ultra pure water system

Llegamos al nivel de pureza del agua adecuado? Esto daña las fibras? Se reduce la actividad de tritio?

3.5 Characterization and R&D on background shields

3.5.1 Passive shield (lead)

3.5.2 Active shield (cosmic veto)

Chapter 4

Tritium monitor prototypes

4.1 Preliminary prototypes, TRITIUM-IFIC 0, TRITIUM-IFIC 1 and TRITIUM-Aveiro

4.1.1 Tritium-IFIC 0

4.1.2 Tritium-IFIC 1

4.1.3 Tritium-Aveiro

4.2 Advanced prototype, Tritium-IFIC 2

4.3 Modular TRITIUM prototype for in-situ tritium monitoring

Chapter 5

Simulations

Chapter 6

TRITIUM Monitor results and discussion

6.1 Results from laboratory measurements

6.2 Results from measurements at Arrocampo dam

6.3 Results in simulations

Chapter 7

Conclusions and prospects

Que cosas se han conseguido en este experimento? -> Decir tanto lo que se ha conseguido con el detector como con las investigaciones de componentes del detector (capítulo 3)

Responder a las grandes preguntas: - Podemos medir tritio? Lo podemos hacer en tiempo quasi real? Lo podemos hacer a la actividad que queríamos? - Que sensibilidad se ha llegado a conseguir? - Estabilidad temporal? - Precio? - Comparación con respecto al resto de experimentos - Efecto del shield - Medidas a varias actividades

Appendices

Appendix A

Birks coefficient study

Aún faltan cosas por decir.

Appendix B

More things

Aún faltan cosas por decir.

Appendix C

And even more things

Y más cosas aún.

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