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Characterisation of a ⁹⁰Sr based electron monochromator

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Abstract

This note describes the characterisation of an energy filtered ⁹⁰Sr source to be used in laboratory studies that require Minimum Ionising Particles (MIP) with a kinetic energy of up to ~2 MeV. The energy calibration was performed with a LYSO scintillation crystal read out by a digital Silicon Photomultiplier (dSiPM). The LYSO/dSiPM set-up was pre-calibrated using a ²²Na source. After introducing the motivation behind the usage of such a device, this note presents the principle and design of the electron monochromator as well as its energy and momentum characterisation.

1. Introduction

In the context of calorimetry R&D for future linear collider experiments, several technologies are currently under study within the CALICE [1] collaboration, all of them aiming at unprecedented highly granular segmentations to be used with particle flow algorithms [2].

In particular, an electromagnetic calorimeter based on small plastic scintillators coupled to silicon photomultipliers (SiPM) has been proposed. A study has been started at CERN on the characterisation of the light yield and its uniformity of small plastic scintillators, of sizes ranging from $10 \times 10 \times 2$ mm³ to $30 \times 30 \times 5$ mm³. These characterisations are performed using various light containment solutions, such as wrapping the scintillator with an Enhanced Specular Reflective¹ (ESR) foil or painting its faces with a reflector paint².

To produce the scintillation light in such laboratory set-ups, electron sources are generally used, in particular the β emitting ⁹⁰Sr isotope, which in combination with the decay of the daughter nucleus Y-90 provides a continuous electron energy spectrum extending up to ~ 2.2 MeV, as shown in Fig. 1.

In order to properly quantify the yield and uniformity of the scintillator under test, it is important to have knowledge of the amount of energy deposited by the traversing electrons. This can be achieved by the use of Minimum Ionising Particles (MIPs). Fig. 2 (top) shows the stopping power (dE/dx) of electrons in a polyvinyltuolene-based plastic scintillator. The lower plot shows the corresponding range. The density of plastic scintillator is 1.03 g/cm^3 . Both plots were generated by the ESTAR program provided by NIST³

Electrons above ~ 1 MeVcan be considered MIPs and will traverse scintillating tiles of few mm thickness as under study for the ECAL application. Therefore, the goal is to prevent elec-

¹3M ESR

³NIST ESTAR http://physics.nist.gov/PhysRefData/Star/Text/ESTAR.html



Figure 1: β spectrum of ⁹⁰Sr and the daughter nucleus ⁹⁰Y.

²Saint-Gobain BC-620



Figure 2: Top: Stopping power (dE/dx) of electrons in a polyvinyltuolene-based plastic scintillator. Bottom: Range of electrons in the same material. The x-axis corresponds to the kinetic energy of the electrons.

trons of lower energy from reaching the scintillator under test, which can be done by selecting the electron momentum using a magnetic field.

The following section briefly describes how such a solution is implemented in the form of a compact electron monochromator.

2. The electron monochromator

The device described here is based on the original design by the *Institut de Recherches Subatomiques*, Strasbourg, France. From their technical drawings, shown in Appendix B, the manufacturing of the tungstene pieces was externally contracted, all other components were fabricated at CERN. The ⁹⁰Sr source itself was inserted by CERN's radio-protection group.

The electron monochromator, illustrated on Fig. 3 consists of three main parts. The body is made of tungsten which ensures efficient shielding in all directions. The source housing, which contains the \sim 350 MBq ⁹⁰Sr source, is also made of tungsten and is permanently bolted to the side of the main body. A collimated hole is present inside the source housing pointing inwards, allowing the electrons to escape. The solenoid coil, mounted on top of the main body, is made of 16 layers of 0.80 mm coated copper wires, arranged in 56 windings per layer. Moreover, the coil is equipped with an ARMCO soft iron core to enhance the generated magnetic field. A cavity inside the main body allows the electrons to travel from the source casing and, in the presence of a magnetic field, bend towards the exit hole on the bottom side of the body. The magnetic field is guided to the cavity via soft iron bars on the sides and additional soft iron cylinders on the level of the cavity.

When no current is applied to the solenoid, the electrons exiting the source casing are stopped inside the tungsten structure, allowing for safe operation and handling of the device. A small Hall probe allowed to measure the magnetic field in between the two soft iron cylinders as a function of the magnet current. The result is shown in the upper plot of Fig. 4. Dismantling and re-assembly of the source (with the 90 Sr - source removed) showed that the current-field relation changed. This is due to the fact that the width of the air gaps between the various soft iron pieces depend on the applied torque to the screws. For relatively high fields (several thousand Gauss) the air gaps lead to losses. It was however demonstrated that the relation between relative rate and actual magnetic field, as shown in the lower plot of Fig. 4, is not affected.



Figure 3: Technical drawings of the electron monochromator. The solenoid coil and the 90 Sr tungsten casing can be seen. When a current is applied to the coil, the electrons are deviated to the collimated opening on the bottom. Overall dimensions: $130 \times 100 \times 60$ mm.

Proper use of the device requires to know the relation between the solenoid current (or the magnetic field in between the soft iron cylinders) and the electron momentum, as well as the spread of the selected momenta. The following sections describe how the electron monochromator was characterised using a LYSO crystal coupled to a dSiPM.



Figure 4: Top: The magnetic field in between the two soft iron cylinders is plotted versus the applied magnet current. Bottom: The relative intensity at the output of the monochromator is plotted versus the magnetic field. The measurement has been performed with plastic scintillators. The counts are collected in typically 100 s.

3. Experimental set-up

The AX-PET collaboration [3] has developed a PET demonstrator [4] based on axially arranged layers of scintillator crystals read out with conventional analogue silicon photomultipliers (SiPM). In a second phase, an alternative readout has been studied [5] using digital silicon photomultipliers (dSiPM) [6] developed by Philips.

One of these dSiPM arrays, coupled to a LYSO crystal, has been used to characterise the electron monochromator. The dSiPM array used has a surface $32 \times 32 \text{ mm}^2$, and its type is DPC-6400-22-44. It consists of 16 *dies* (7.15 × 7.9 mm²), each segmented into 4 *pixels*. Each $3.2 \times 3.9 \text{ mm}^2$ pixel is comprised of a matrix of 6400 avalanche photo-diodes operated in Geiger

mode, each capable of detecting single photons. Details on the operation of the dSiPM can be found in [5].

The LYSO crystal, $3 \times 3 \times 3$ mm³ in size, is coated with a TiO₂ reflective paint on 5 of its sides in order to improve light containment. The bare face is glued to one of the dSiPM pixels, effectively covering ~ 72% of its surface. Therefore, only ~ 4600 of the 6400 cells are exposed to the light of the crystal.

Before characterising the electron monochromator, a calibration of the non-linear response of the LYSO/dSiPM arrangement is required. This is performed by comparing the dSiPM amplitude, *i.e.* the number of dSiPM cells fired, to the known energies of the photons emitted by a 22 Na source. The experimental set-up for this non-linearity measurement, sketched in Fig. 5, consists of the following elements:

- a ²²Na source for the dSiPM + LYSO calibration
- the dSiPM array and its readout electronics
- the LYSO crystal, glued to the dSiPM array
- a laptop for the dSiPM readout



Figure 5: Schematic representation of the experimental set-up for measuring the non-linearity of the LYSO/dSiPM arrangement.

The results of the calibration are shown in the following section.

4. dSiPM calibration

The number of dSiPM cells fired for a given energy deposit E in the LYSO crystal can be written as follows:

$$N_{cells} = \alpha \cdot \left[1 - \exp\left(-\frac{E}{\beta}\right) \right] \tag{1}$$

where α and $p\beta$ are calibration constants. This functional relation reflects the signal saturation imposed by the combinatorics due to the finite number of dSiPM cells (4600). A cell which is hit by two and more photons is still only counted as 1 hit. To determine the calibration constants, the LYSO was exposed to a ²²Na source as shown on Fig. 5 and a dSiPM response spectrum was acquired; it is shown in Fig. 6 (left). The two main peaks correspond to the positronium annihilation, resulting in two photons of 511 keV, and a nuclear transition line at 1270 keV, respectively. They were both fitted with a Gaussian distribution and their mean values, together with the origin (0,0) appear on the graph of Fig. 6 (right). This graph was fitted with the function shown on Eq. 1 and the parameters α and β were obtained.



Figure 6: Left: Spectrum of the number of dSiPM cells fired under the ²²Na source, showing the 511 keV positronium annihilation peak, the 1.27 MeV nuclear transition peak, and their respective Compton edges. Right: Calibration fit using the identified peaks, including the point corresponding to (E = 0 keV; $N_{cells} = 0$).

Using the relation between the numbers of dSiPM cells fired and the energy deposited in the LYSO crystal, it is now possible to characterise the electron monochromator in energy and momentum, as detailed in the next section.

5. Electron monochromator characterisation

The experimental set-up, sketched in Fig. 7, is similar to the calibration with the ²²Na source, however now the ⁹⁰Sr β source is being used. The electron monochromator exit hole was placed above the LYSO crystal, as close as possible. The procedure was to set various solenoid currents and for each, acquire a dSiPM response spectrum.

The number of fired dSiPM cells, N_{cells} , can be converted into an energy deposit using the previously obtained calibration:

$$E = -\beta \cdot \ln\left(1 - \frac{N_{cells}}{\alpha}\right) \tag{2}$$



Figure 7: Schematic representation of the experimental set-up for the energy calibration of the electron monochromator.

and the momentum p can be calculated as follows:

$$p = \sqrt{(E+m_e)^2 - m_e^2}$$
(3)

where $m_e = 511$ keV is the electron mass.

Fig. 8 shows an example momentum spectrum obtained at a solenoid current of 1.2 A. Distributions for other current setting can be found in Appendix A. The main peak corresponds the electrons passing the monochromator. This peak moves linearly with the settings of the solenoid current. The background does not depend on the magnetic field inside the monochromator and is attributed to bremsstrahlung photons (soft X-rays) that are produced by electrons inside the steel/tungsten casing. While these X-ray photons are visible in the high-Z LYSO crystals, the Z^{4-5} dependence of the photoelectric effect would suppress their detection in a plastic scintillator.

Table 1 summarises the results of the full measurement campaign, showing the observed number of dSiPM cells fired, the calculated energies, momenta, and respective resolutions for each setting of the solenoid current. Below 0.4 A, no peak was visible above the bremsstrahlung background. Beyond 1.6 A, the end of the ⁹⁰Sr β -emission spectrum is being approached, and the rate decreases rapidly. The precision of the momentum selection is generally better than 10%, for momenta above 1.5 MeV/c it approaches 5%.

In order to verify the expected linearity between the electron momentum and the solenoid current, each value was added to a graph fitted with a first degree polynomial, as shown in Fig. 9. A good linearity is observed over the current range explored. Finally, Fig. 10 shows the energy and momentum selection resolutions as a function of the solenoid current. We observe that a momentum selection resolution of $\sim 5 \%$ can be achieved for currents above 1.4 A. This solenoid current value corresponds to our working point for MIP selection in our 2 mm thick scintillator studies.

It should be noted that, as described above, the relation between the magnet current and the



Figure 8: Example of a momentum distribution at a solenoid current of 1.2 A. The red curve is a Gaussian fit that corresponds to the selected electrons. The background corresponds to bremsstrahlung photons escaping the monochromator casing, and is independent of the solenoid current.

selected momentum, may change due to external effects, e.g. if the torque of the screws are modified. For typical applications, the set-up may be used selecting the magnet current which provides the highest rate at the exit of the monochromator. This corresponds to electrons with momenta of about 1.4 - 1.6 MeV/c, i.e. $E \approx 1$ MeV. According to Fig. 2 such electrons can be considered as minimum ionising particles.

6. Conclusion

An electron monochromator, based on a high-activity ⁹⁰Sr source, has been manufactured, assembled, and characterised by using a LYSO crystal coupled to a digital silicon photomultiplier array. After determining the non-linearity of the dSiPM/LYSO response with a ²²Na source, a

<i>I</i> [A]	N _{cells}	E [keV]	σ_E [keV]	σ_E/E [%]	p [keV/c]	σ_p [keV/c]	σ_p/p [%]
0.40	1018	248	83	33.5	561	108	19.3
0.60	1839	488	79	16.2	858	90	10.5
0.65	1943	522	79	15.1	898	88	9.8
0.80	2591	760	81	10.7	1164	87	7.5
1.00	3243	1050	88	8.4	1474	92	6.2
1.10	3504	1187	94	8.0	1619	97	6.0
1.15	3594	1237	91	7.4	1672	94	5.6
1.20	3750	1328	96	7.2	1767	100	5.7
1.30	3986	1485	103	6.9	1929	105	5.4
1.40	4138	1594	106	6.6	2042	111	5.4
1.60	4467	1871	108	5.8	2327	108	4.6

Table 1: Measurement results for various solenoid current settings across the ⁹⁰Sr emission spectrum. E denotes the kinetic energy of the electrons. σ_p corresponds to the width of the Gaussian fit to the peaks for each current. σ_E is calculated by error propagation following eq. 3

study of the electron momentum selection as a function of the solenoid current has been performed. As expected, a linear dependency was observed. Additionally, the resolution of the momentum selection has been measured for various currents. It was shown that a 5 % spread in momentum selection can be achieved for electrons above ~ 2 MeV/c, rendering this device very useful for small laboratory experiments requiring MIPs.

In typical applications, the monochromator will be used in combination with a trigger counter. In the tile scan set-up for the Linear Collider Detector studies, a cross arrangement of two 1 mm thick squared scintillating fibres is mounted in about 1 cm distance from the exit slit. The effective trigger area = 1 mm^2 . The typical trigger rate is of the order 10 Hz.

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Figure 9: Electron momentum as a function of the electron monochromator solenoid current. The red curve corresponds to a linear fit to the data points.

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Figure 10: Energy (left) and momentum (right) resolutions for the various solenoid current settings. A momentum selection of ~ 5 % can be achieved for currents above 1.4 A.

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A. Additional figures

Figure 11: Momentum distributions for various solenoid current values



Figure 12: Momentum distributions for various solenoid current values

B. Technical drawings























