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Propagation Speed of $R_{\gamma-511}$ keV in Plastics

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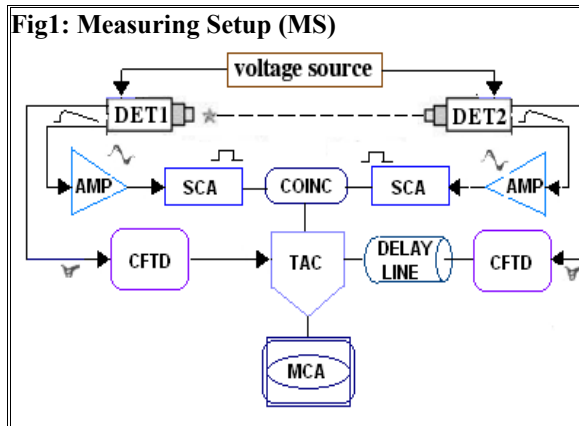
Abstract: fitted central peaks of the coincidence spectra resulting from measuring the propagation speed (PS) of ^{22}Na -isotope γ -radiation 511 keV in plastics

Key Words: propagation speed, ^{22}Na -isotope, electronic coincidence method, plastics

1) Introduction - The propagation speed (PS) of visible light - a small frequency range in the large frame of electromagnetic radiations (ER)- in air was measured, during the last hundred years⁽¹⁾, using a great deal of different methods, with high precision results being achieved.

Further improvements of detection and electronic measuring systems allowed to determine such parameters as PS by using $R\gamma$ ^(2,3,4,5,6,7) with an advantage that such experiments could also be performed in a larger variety of propagation media non-transparent to visible light. In this context, beside our already performed measurements in air⁽⁷⁾, in a good extent comparable to the CODATA value⁽⁸⁾, we extended such measurements in plastics as a propagation media. As already well settled⁽²⁻⁷⁾, to perform such PS measurements the availability of a $R\gamma$ source in which two $R\gamma$ are emitted simultaneously in opposite directions turns out to be essential to the feasibility of the experiment, as far as no reflection techniques could be used. Such suitable source, in all cases, was the positron emitter ^{22}Na placed in a metal container in which the positrons are stopped and annihilated when reacting with the medium electrons, in such way originating -as it is very well established from momentum/energy conservation laws⁽⁹⁾- two $R\gamma$ -511 keV each, both emitted simultaneously in opposite directions.

2) Experimental – The measuring setup [MS (Fig21/photo)] included two $R\gamma$ detectors, DET1



(photomultiplier XP-2020Q + BaF₂ scintillator) and DET2 (photomultiplier XP-2020 + CsF scintillator), each of them connected to an electronic fast-slow coincidence circuit [slow branch: amplifier (AMP), timing single channel (SCA), universal coincidence (COINC); fast branch: constant fraction timing discriminator (CFTD); time to pulse amplitude converter (TAC). Finally, the slow-fast coincidences

were recorded on an analog-digital- converter/multi-channel (MCA). More detailed explanations about construction and performance of such an MS can be found elsewhere^(10,11). The experiments consisted in the measurements of the eventual shifts of the fitted coincidence spectra peaks when different

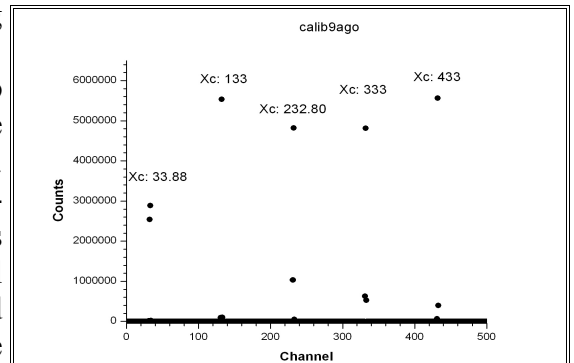


Fig2 - Time-Calibrator output spectrum

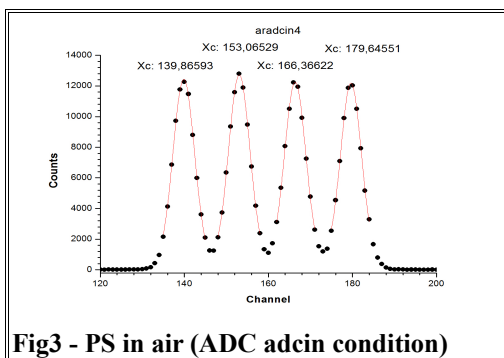
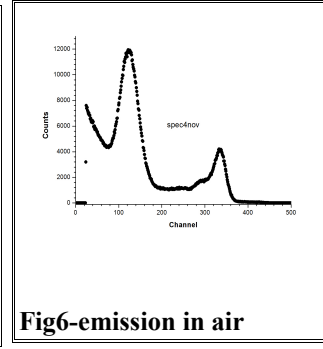
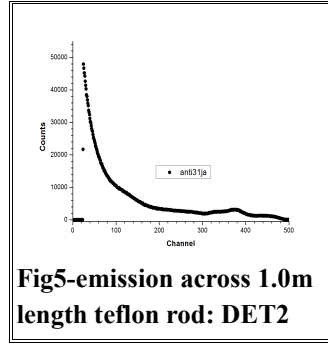
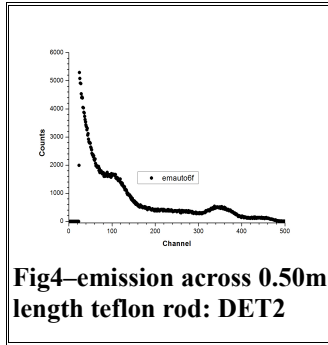


Fig3 - PS in air (ADC adcin condition)

plastic rods were interposed. The two oppositely emitted $R\gamma$ originated from a $\sim 30 \mu\text{Ci}/^{22}\text{Na}$ γ -emitter, as far as they appeared as coincidence spectra displayed in a Multi-Channel (MC).

As a first step of the experiments, it was measured the *time calibration* of the MS by using a Time Calibrator (TC), range $0.08 \mu\text{s}$ -period $0.01 \mu\text{s}$, which produces two pulses with highly precise variable delays between their outputs; which, by their way, were directed to the TAC whose amplitude outputs is related to those delays (Table I). Finally, the average *time calibration*



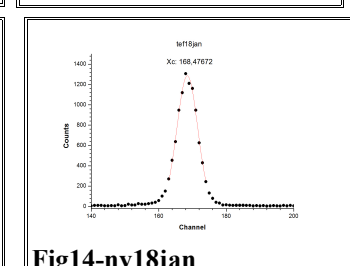
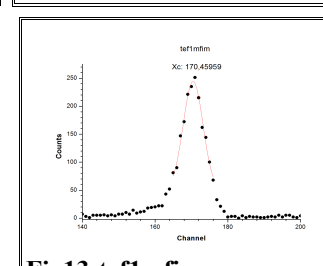
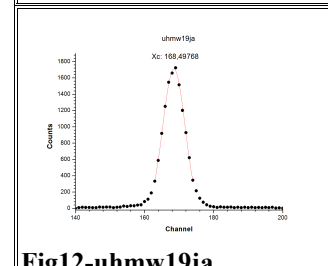
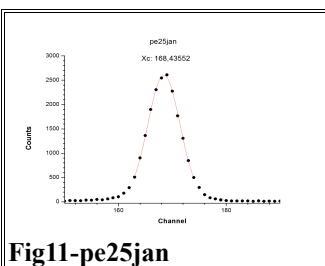
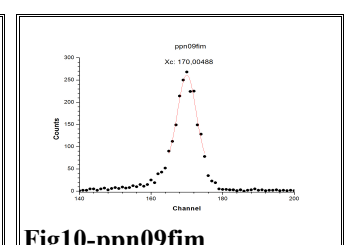
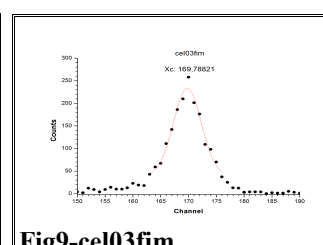
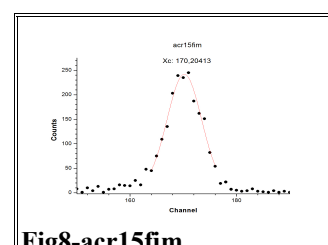
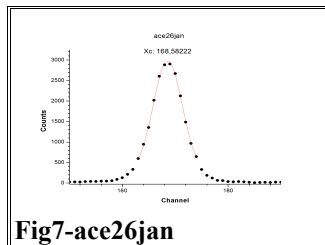
displayed by the MS was **0.1002204 ns/ch**, a result extracted from fitting (Fig2).

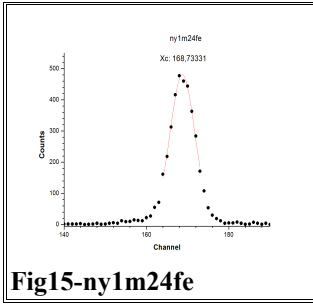
As beginning coincidence experiments, in order to fix standards spectra and time-energy calibration in real measurement conditions of the MS, PS-adcin in air was

performed (Fig3) with an estimation: $PS = 300,998.90 \text{ km/s}$ ($1.20 \text{ m} \rightarrow 179.64551 \text{ ch} - 139.86593 \text{ ch} = 39.77958 \text{ ch} \rightarrow 3.9867254 \text{ ns} - 1.20 \times 10^{-3} \text{ km} / 3.9867254 \times 10^{-9} \text{ s} = 300,998.90 \text{ km/s}$). The following PS experiments were performed with the **Ry-511 keV**, detected by **DET2**, after propagating in **1,0 m** and **0,50 m** plastic rods with **25 mm** diameter; the two experiments with each plastic road in the two multiple double dimensions were performed in order to get the time-standards relative to well known propagating distances. Also, in order to make sure that the **Ry-511 keV** would be still present in **DET2** after it's propagation through the plastic rods, emission spectra are displayed in (Figs4-5): a comparison between the ^{22}Na emission spectra in air and after propagating across the most dense plastic, teflon, used in the following experiments; compared to air (Fig6), it may be realized from the displays that the **Ry-511 keV** is certainly attenuated but, even so, still visible. As the next steps, coincidence experiments were performed and the fitted peak's centers of the measured spectra were determined (Tables I-II). All the measured coincidence spectra were fitted with "gaussian function" (eq.1), as founded in the **ORIGIN** software⁽¹²⁾.

Table I: One-Meter (1 m) Interposed Plastic Rods

interposed 1m plastic rods	archives	central fitted channel
acetal	<i>ace26jan</i>	168.58222 (fig7)
acrílico	<i>acr15fim</i>	170.20413 (fig8)
celeron	<i>cel03fim</i>	169.78821 (fig9)
polipropileno	<i>ppn09fim</i>	170.00488 (fig10)
polietileno	<i>pe25jan</i>	168.43552 (fig11)
uhmw	<i>uhmw19ja</i>	168.49768 (fig12)
teflon	<i>tefl1mfim</i>	170.45959 (fig13)
nylon1	<i>ny18jan</i>	168.47672 (fig14)
nylon2	<i>ny1m24fe</i>	168.73331 (fig15)
air spectra	<i>aradcin4</i>	139.86593-153.06529-166.36622-179.64551
emission – 1m teflon interposed	<i>anti31ja</i>	fig6
emission in air	<i>emadinco</i>	fig7





The full-half-width (**fhwm**) of the so fitted coincidence spectra displayed 7-8 channels, **0.7015428ns-0.8017632ns**, according the **MS** calibration; reasonable high values when compared to the time differences between the coincidence spectra fitted central peaks. Even so only the coincidence spectra fitted central peaks, without error bars or **fhwm**, were considered in the present **PS** estimations.

Table II: Half-Meter (0,50 m) Interposed Plastic Rods

interposed 0.50 m plastic rods	archives	central fitted channel
acetal	<i>aceadc21</i>	150.54677 (fig16)
acrílico	<i>acradc12</i>	150.65962 (fig17)
celeron	<i>cel10adc</i>	150.63215 (fig18)
polipropileno	<i>ppnadc13</i>	150.11976 (fig19)
polietileno	<i>penadc14</i>	150.28886 (fig20)
uhmw	<i>uhmwadin</i>	150.09526 (fig21)
teflon	<i>tefadcin</i>	150.60433 (fig22)
nylon	<i>nyfadcin</i>	150.60433 (fig23)
emission – 0,50m teflon interposed	<i>emauto6f</i>	fig6

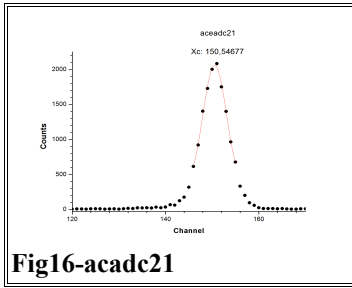


Fig16-acadc21

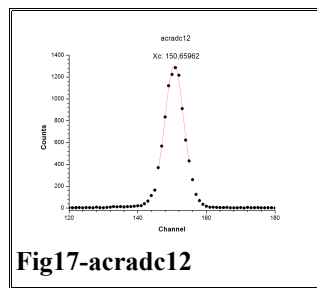


Fig17-acradc12

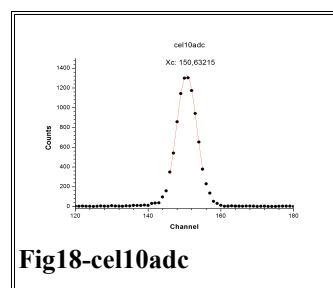


Fig18-cel10adc

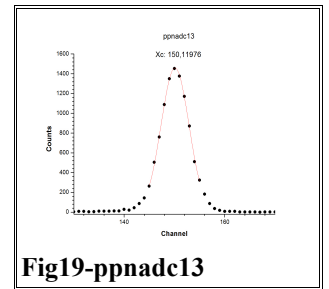


Fig19-ppnadc13

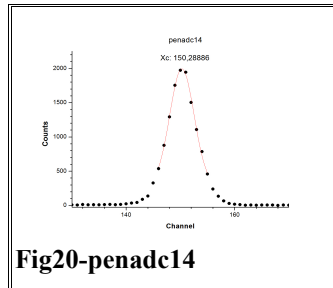


Fig20-penadc14

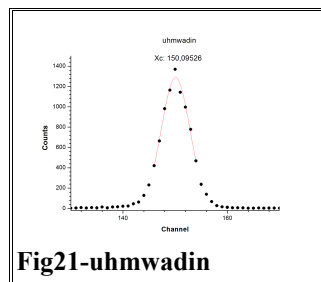


Fig21-uhmwadin

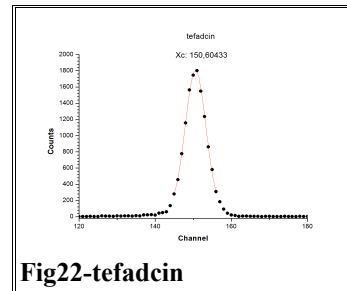


Fig22-tefadcin

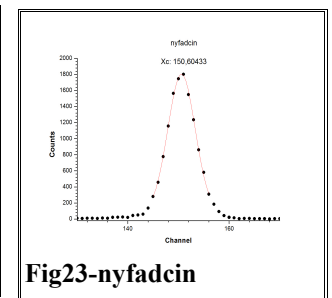


Fig23-nyfadcin

Table III: Time Channel Relationships – PS Parameters

Plastic Rods Interposed	Time/Channels Relations ($\Delta t/\text{ch}$)– 0,50m Plastics	Propagation Speed – PS (km/s)
polipropileno ($\delta= 0.91$)	170.00488 – 150.11976= 19.88512 ch \rightarrow 1.9928946 ns	250,891.34
uhmw ($\delta= 0.93$)	168.49768 – 150.09526= 18.40242 ch \rightarrow 1.8442978 ns	271,105.89
polietileno ($\delta= 0.95$)	168.43552 – 150.28886= 18.14666 ch \rightarrow 1.8186655 ns	274,926.86
nylon1 ($\delta= 1.13$)	168.47672 – 150.60433= 17.87239 ch \rightarrow 1.791178 ns	279,145.90
nylon2	168.73331 – 150.60433= 18.12898 ch \rightarrow 1.8168936 ns	275,194.98
acrílico ($\delta= 1.25$)	170.20413 – 150.65962= 19.54451 ch \rightarrow 1.9587586 ns	255,263.71
acetal ($\delta= 1.42$)	168.58222 – 150.54677= 18.03553 ch \rightarrow 1.807528 ns	276,620.88
celeron ($\delta= 1.45$)	169.78821 – 150.63215= 19.15606 ch \rightarrow 1.9198279 ns	260,440.01
teflon ($\delta= 2.2$)	170.45959 – 150.60433= 19.85526 ch \rightarrow 1.989902 ns	251,268.65

3) Concluding Remarks –

- a) the electronic γ - γ coincidence method showed to be a valuable method to measure **PS** of electromagnetic radiation in plastics, even when measured in very short distances.
- b) in the above estimations it was taken into account only the **fitted peak's centers** of the coincidence spectra, neglecting any deviation concerned these values as done, for instance, by the **fwhm** of those spectra.
- c) the **PS** final results are closely related to the **MS** calibration conditions due to crystal scintillator's shapes and sizes, as well as detectors distances to emitting source.
- d) at a first glance to the refractive indexes it would be difficult to associate **PS** changes to the materials densities in a regular way; different from what expected, lower densities may be related to lower **PS**
- e) going on with the possibility of measuring such **PS** in material media non-transparent to visible light, a topic that the **LCA/CBPF** is presently extending to metals.

4) References -

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- (11) "Radiation Detection and Measurements" - Glenn F. Knoll; J. Wiley&Sons, eds. 1979-1989.

(12) www.originlab.com – equation parameters:

y = maximum height

y_0 = initial height

A = amplitude

σ = average half-width

μ = central channel



Fig24 - MS photo: LCA/CBPF

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