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Absolute efficiency calibration of a coaxial HPGe detector for quantitative PGAA and T-PGAA

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Abstract. Following international guidelines by the IEEE and a well-established calibration
procedure developed at the Hungarian Institute of Isotopes in Budapest, the experimental
absolute efficiency calibration of an Ortec n-type coaxial high-purity germanium (HPGe)
detector (GMX Profile series) has been performed, in view of direct application to quantitative
PGAA and T-PGAA investigations of Cultural Heritage multicomponent manufactures.

1. Introduction
Among the several characteristics of a γ-ray detector, the absolute full-energy-peak (FEP)
efficiency (i.e., the ratio of the number of counts under the photopeak to the total number
of γ-rays emitted by the source) plays a vital role for quantitative spectroscopy, especially when
complex γ-ray spectra involving many lines are under scrutiny. In particular, the development
of Prompt-Gamma Activation Analysis (PGAA) and Time-resolved Prompt-Gamma Activation
Analysis (T-PGAA) capabilities at pulsed neutron sources has allowed to expand the potential
of PGAA at continuous sources for the study of functional materials and Cultural-Heritage
samples [1–3]. These investigations require the use of high-resolution γ-ray detectors to resolve
the complex spectra of multicomponent samples.

In this field, semiconductor detectors are much appreciated in comparison to inorganic
scintillators such as NaI(Tl), due to their superior energy resolution when operated at low
temperatures. However, this improved energy resolution comes along with a major drawback,
since the generally smaller sizes of common germanium detectors, together with the lower atomic
number of Ge with respect to iodine, lowers the FEP efficiency ($\epsilon_{fep}$) by an order of magnitude
in typical cases [4].

For this reason, the assessment of the absolute $\epsilon_{fep}(E)$ curve (i.e., $\epsilon_{fep}$ as a function of
photon energy) of a HPGe detector becomes a crucial analytic passage, also taking into account
the unknown exact functional dependence of efficiency on energy (only partially fixed by the
Table 1: Source features, from source datasheet and LNE-LNHB/CEA nuclear-data tables [9]. The most important acquisition parameters are reported as well: $A_0 =$ reference activity (3% uncertainty with 95% confidence), $t_1 = $ half time, $t_{live} = $ livetime, i.e., total time of actual count acquisition with no dead time included, and $d = $ source-detector distance.

|       | $A_0$                     | $t_1$     | $t_{live}$ | $d$ | $t_{live}$ | $d$ |
|-------|---------------------------|-----------|------------|-----|------------|-----|
| $^{60}$Co | $(42.10 \pm 1.26) \text{ kBq}$ | $5.2711 \pm 0.0008 \text{ y}$ | 7200 s | 11.5 cm | 300 s | 11.5 cm |
| $^{133}$Ba | $(36.60 \pm 1.10) \text{ kBq}$ | $10.539 \pm 0.006 \text{ y}$ | 7200 s | 11.5 cm | 300 s | 11.5 cm |
| $^{137}$Cs  | $(41.10 \pm 1.23) \text{ kBq}$ | $30.05 \pm 0.08 \text{ y}$ | 7200 s | - | 1 cm | - |
| $^{241}$Am  | $(39.60 \pm 1.19) \text{ kBq}$ | $432.6 \pm 0.6 \text{ y}$ | 7200 s | 11.5 cm | 300 s | 11.5 cm |

application of empirical or semi-empirical models [5]), as well as limits to the overall accuracy due to unavoidable sources of uncertainty hidden in the geometry of the experimental setup.

The 1996 IEEE protocol, which establishes terminology and standard test procedures for germanium $\gamma$-ray detectors [6], contains popular guidelines to perform various types of calibrations. Its integration to the novel efficiency-calibration procedure developed at the Hungarian Institute of Isotopes in Budapest [5,7] allows for the improvement of some practices, such as the assessment of the system non-linearity, which is mainly due to the analog-to-digital converter (ADC), and is cause to major FEP distortions that cannot be neglected when (T-)PGAA is performed because of the very low amplifier gains employed in this technique [8]. Starting from a careful mix of such protocols, the experimenter is allowed to perform the $\epsilon_{fep}$ calibration of a HPGe detector over a wide energy range ($\sim 0.05-11.00 \text{ MeV}$), typically making use of a large set of calibrated and noncalibrated samples of radionuclides over the low-energy region (up to about 2 MeV), while exploiting cascades from thermal-neutron capture on non-calibrated, non-commercial light-nucleus targets (such as $^{14}$N or $^{35}$Cl) over the high-energy counterpart (above 2 MeV) [5].

In the present work, a practical application of this procedure is presented for the assessment, over the low-energy region, of the absolute $\epsilon_{fep}(E)$ curve of an Ortec n-type coaxial HPGe detector (GMX Profile series).

2. Experimental setup
The thermostated HPGe detector shown in Fig. 1 was employed to collect a double set of $\gamma$ spectra of an ensemble of 4 calibrated radionuclides (reference time: $t_0 = 01/11/2005$) in a shielded area located in the Target Station 2 (TS2) building at the ISIS pulsed neutron and muon source [10], UK. Measurements included “beam-off” background (both measurement sets were collected in the course of a maintenance shutdown of all ISIS beamlines, in order to minimize unwanted signals in the spectra).

A summary of source features and major acquisition parameters is reported in Table 1. In step with the IEEE protocol [6], discoidal standards were used with an active nucleus of $\leq 2 \text{ mm}$ in diameter, and the livetime of the background collection was consistent with the one of any source spectrum. Distances from the crystal - which do not include the gap (2.6 cm) between the endcap and crystal - were shortened in comparison to the customary 25-cm value, due to the very low activities of the sources. These distances were chosen as a reasonable trade-off.
between maximization of signal-to-noise (S/N) ratio and minimization of coincidence summing (which induces FEP loss). Additional distances for $^{137}$Cs were taken for future employment in simulations.

Following the same route, a second batch of measurements (Set 2) was taken in order to correct source placement (according to a coaxial geometry, Fig. 2) and fix the non-optimal polarization of the semiconductor junction by switching to -3800 V in the high-voltage (HV), compared to the initial -1800 V value.

No collimation and/or Compton-suppression arrangement were used.

3. Data reduction and analysis

After a channel-by-channel background subtraction from any source spectrum, corresponding spectra in Sets 1 and 2 were compared to one another. Though the former show larger S/N ratios (a neat example is shown in Fig. 3 for barium), the latter were chosen for subsequent analysis due to the significant reduction in the left skewing of peaks following HV correction (an example is presented in Fig. 4 for cobalt), with consequent enhancement of spectral resolution, lowering of the low-energy threshold down to about 20 keV, and removal of any possible multi-peak fitting of single lines.

The first part of the data analysis was performed by Hypermet-PC software [8,11], which is currently considered the most reliable - though closed-source - package for the automated analysis of complex $\gamma$-ray spectra [12]. The procedure outlined below follows the indications reported in [5]. After direct editing and update of the Hypermet-PC database of nuclear data (originally dating back to the early 2000s), the following relationship

$$E = a + m \cdot Ch,$$

where $a$ (in keV, like $E$) and $m$ represent the intercept and slope, respectively, of the calibration line, while $Ch$ is the position of the peak centroid (in channels), was used to perform an initial energy pair-calibration (i.e., the conversion of channels into energies) for each spectrum with no consideration - at a first approximation - of non-linearity effects introduced by the ADC.

This calibration was used to perform a first fit of each spectrum (“peaklist 0”), taking into account that the $i^{th}$ peak profile in the $j^{th}$ channel is formalized as:
\[ p(j) = \Gamma_i e^{-\frac{(j-j_{i,0})^2}{2\delta^2}} + \frac{1}{2} A_i e^{-\frac{\delta}{\beta} \text{erfc} \left( \frac{\delta}{2\beta} + \frac{j-j_{i,0}}{\delta} \right)}, \] (2)

where the first term in the right part of the equation is a Gaussian of amplitude $\Gamma_i$ and variance $\frac{\delta}{\sqrt{2}}$, and the second one is an exponentially-modified Gaussian (EMG) - of amplitude $A_i$, variance $\frac{\delta}{\sqrt{2}}$, and exponential-decay time constant $\beta$ - representing the left skewing.

Figure 2: Source placement for Set 2.

An additional baseline correction of the form:

\[ S_i \text{erfc} \left( \frac{j-j_{i,0}}{\delta} \right) + \frac{1}{2} W_i e^{-\frac{\delta}{2\tau} \text{erfc} \left( \frac{\delta}{2\tau} + \frac{j-j_{i,0}}{\delta} \right)} + c_0 + c_1 j + c_2 j^2, \] (3)

was applied, including in progressive order:

- an error function of amplitude $S_i$, which accounts for photons scattered at small angles;
- a “tail” contribution (EMG with exponential-decay time constant $\tau \gg \beta$), which accounts for detector surface-effects, especially at very low energies;
- a parabolic contribution with polynomial coefficients $c_0$, $c_1$, and $c_2$.

A higher-order refinement of the above fit (“peaklist 1”) was, then, performed by a non-linearity assessment accomplished by construction of a non-linearity curve via orthonormal-polynomial least-square (OPLS) fit of source peak-positions in peaklist 0. This procedure - which consists of measuring the positions of the FEP centroids in the spectrum of a multi-emitter, and then evaluating their displacement from a calibration linear curve [5,8] - is more accurate than the one suggested by the IEEE, which implies the use of a high-precision pulse generator at the test-input terminal of the pre-amplifier to “simulate” the electronic-noise contribution to the FEP FWHM [6].

Being only one multi-emitter ($^{133}$Ba) available in the calibration set, this initially limited result accuracy over the energy range $\geq 400$ keV. The problem was bypassed via an ad hoc correction of centroid errors for $^{137}$Cs and $^{60}$Co by picking appropriate values reported in LNE-LNHB/CEA tables.

The ensemble of all peaklists 1 was finally used to perform the assessment of the intrinsic part of $\epsilon_{fep}(E)$, namely $\epsilon_{int}(E)$, that can be defined as [5]:

\[ \epsilon_{int} = \frac{C}{P_\gamma N_{\text{live}}^{\text{abs calibrated source}}} \] (4)
Figure 3: Top: $^{133}$Ba $\gamma$ spectrum (Set 1) up to energies of the order of 0.5 MeV. Gamma emissions from LNE-LNHB/CEA tables: 1 = 53.1622(18) keV, 2 = 79.6142(19) keV, 3 = 80.9979(11) keV, 4 = 160.6121(16) keV, 5 = 223.237(2) keV, 6 = 276.3992(21) keV, 7 = 302.8512(16) keV, 8 = 356.0134(17) keV, and 9 = 383.8491(12) keV. Visible random sums: RS 1 = peak 3 + K$_{\alpha,\beta}$ X lines of Cs, RS 2 = peak 7 + K$_{\alpha,\beta}$ X lines of Cs, RS 3 = peak 8 + K$_{\alpha,\beta}$ X lines of Cs, and RS 4 = peaks 2-3 + peak 8. Bottom: same spectrum (zoom) up to energies of 1 MeV, with a tentative assignment of the annihilation peak at 510.999 keV.
Figure 4: Top: 1.173- e 1.332-MeV lines of $^{60}$Co in the Set-1 spectrum. Bottom: same lines in the correspondent Set-2 spectrum.
Table 2: Coefficients recovered by OPLS fitting (ROOT software) of discrete efficiency points associated with radionuclide peaks.

| Grade | $a_0$ ± $a_1$ | $a_2$ ± $a_3$ |
|-------|---------------|---------------|
| 2     | $-1.979 ± 0.216$ | $4.128 ± 0.082$ | $-0.378 ± 0.007$ |
| 3     | $-29.776 ± 0.046$ | $19.347 ± 0.014$ | $-3.074 ± 0.002$ | $0.1556 ± 0.0002$ |

\[ \epsilon_{int} = \frac{C}{P_{\gamma}} \quad (\text{non calibrated source}), \quad (5) \]

where

- $C =$ number of counts for each peak
- $P_{\gamma} =$ $\gamma$-emission probability
- $N(t) = \frac{A_0}{\lambda} e^{-\lambda(t-t_0)} (1 - e^{-\lambda t_{acq}}) =$ number of decays from the start of measurement
- $t_{live} =$ livetime
- $t_{acq} =$ acquisition time.

In this way, discrete efficiencies were calculated for each line of each radionuclide, selecting one source ($^{133}$Ba) as absolutely calibrated, and using it as a normalizing factor for the remaining standards (which practically corresponds, in the log-log scale, to an efficiency shift of $^{60}$Co, $^{137}$Cs, and $^{241}$Am along the vertical axis to match $^{133}$Ba values).

Then, an OPLS fit of the above discrete efficiency points, based on the commonly-used function [13,14]

\[ \ln [\epsilon_{int}(E)] = \sum_{i=0}^{n} a_i [\ln(E)]^i, \quad (6) \]

(linearized $\chi^2$ method) was performed in order to get the desired $\epsilon_{int}(E)$ curve. Due to Hypermet-PC being proprietary software, the obtained curve could not be exported to a regular ASCII file. An external reconstruction of $\epsilon_{int}(E)$ was attempted, starting from fit coefficients $a_i$ calculated by Hypermet-PC. Yet, even taking into account the remapping of energies onto the ($-1,1$) range operated by the software in order to improve the stability of the fitting algorithm [5,15], no adequate correspondence was found, probably due to an unknown compression/expansion factor applied by Hypermet-PC to the curve. Consequently, a parallel fit (again by Eq. 6) of the same discrete efficiency points was performed via ROOT software [16], recovering the results summarized in Table 2 and Fig. 5. Low-grade polynomial functions were used in order to avoid any possible artificial oscillation of the resulting efficiency response curve in those regions covered by sporadic experimental points.

Finally, in Fig. 6 the grade-3 fit for $\epsilon_{int}(E)$ is shown together with its uncertainty region, after normalization to the mean value of the discrete intrinsic efficiency of the barium line at 160.6121 keV.

In order to get the global absolute FEP efficiency $\epsilon_{fep}(E) = \epsilon_{geom} \epsilon_{int}(E)$ for a point-source geometry, a multiplication must be performed by a geometric factor $\epsilon_{geom} = \frac{\Omega}{4\pi}$, with $\Omega = 2\pi \left(1 - \frac{d}{\sqrt{d^2 + r^2}}\right) =$ solid angle “seen” by the detector, $d =$ source distance from the crystal, and $r =$ crystal radius. In our case, at $d = (11.5 + 2.6)$ cm and $r = 2.98$ cm, it is $\epsilon_{geom} = 0.011$, which explains the initial need for reducing the source-detector distance from 25 to 14.1 cm.
Figure 5: Top: lin-lin representation of the absolute intrinsic efficiency curves (grade 2 and 3) obtained by OPLS fitting (ROOT software) of discrete efficiency points associated with radionuclide peaks. Bottom: same curves in a log-log representation.
4. Discussion
A quick look at Table 2 and Fig. 5 suggests that, in spite of the grade-3 fit showing an unphysical growth of the intrinsic efficiency curve in the high-energy region (where absorption cross-sections tend to decrease [7]), this same curve fits best the low-energy portion of the spectrum.

Consequently, we have opted for saving this grade-3 low-energy “patch”, with the purpose to link it - by an appropriate continuity condition imposed on $\epsilon_{int}(E)$ at the matching point - to a high-energy complement obtained by future measurements on other non-calibrated sources, and/or ongoing Monte Carlo simulations of the detector efficiency, and/or similar data from the scientific literature.

5. Conclusions
The absolute efficiency of an Ortec n-type coaxial HPGe detector, used for PGAA and T-PGAA measurements on functional and Cultural Heritage samples at the ISIS pulsed neutron and muon source [2, 3, 17, 18], was determined by the exploitation of a set of calibrated $\gamma$-emitting sources following the procedures outlined in Refs. [5, 6]. The results are in agreement with the values reported in the literature [5, 19, 20].

This work will allow to carry out quantitative T-PGAA measurements on multicomponent samples, primarily addressing functional materials and Cultural Heritage artifacts, as well as quantitative estimates of the absolute $\gamma$ background at ISIS beamlines.

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