L09: In-Situ Contamination & NORMs Measurement

Branislav Stribrnsky Slovak University of Technology Faculty of Electrical Engineering and Information Technology Institue of Nuclear and Physical Engineering

Introduction

- Gamma-ray spectrometry measurements performed on site
 - In-situ Gamma Spectrometry
 - Soil Geometry Mesurement
 - Infinite Plane Geometry Mesurement
 - Ground measurement
 - Ground gammaspectrometry
 - Field gammaspectrometry
- Field of Use
 - Geology
 - Nuclear energy
 - Radiation protection
 - Radiological mapping







History

• 1972

Beck, H. L., DeCampo, J., and Gogolak, C. In Situ Ge(Li) And Nai(Tl) Gamma-Ray Spectrometry.. United States: N. p., 1972. Web. doi:10.2172/4599415.





HASL-258

IN SITU Ge(Li) AND NaI(TI) GAMMA-RAY SPECTROMETRY

September 1972

Health and Safety Laboratory (AEC) New York, New York



History

• 1988

Helfer I. K., Miller K. M. Calibration factors for Ge detectors used for field spectrometry. Health Phys. 1988 Jul;55(1):15-29. doi: 10.1097/00004032-198807000-00002. PMID: 3391774.

	Table 1. ϕ – Unsc	attered flux (y ci	m ⁻² s ⁻¹) at 1 m	above gr	ound per unit	source stre	ngth in the soi	1.*						
Source				(a)	0) - cm ² §	-1								
energy (keV)	0 (Uniform)	0.0625	0.206	0.3	12 0.6	525	6.25	(plane)						
50	1.4403	0.0816	0.2245	0.			Tai	ole 3. Ang	ular corre	ction fact	or (N_c/N_c)).*		
100	2.7744	0.1458	0.3627	ο.										
150	3.3264	0.1702	0.4103	ο.										
200	3.9056	0.1843	0.4550	0.						L/D				
250	4.0640	0.2008	0.4697	0	Energy									
364	4.7184	0.2268	0.5158	0	(Mev)	0.5	0.6	0.7	0.8	0.9	1.0	1.1	1.2	1.3
500	5.3904	0.2519	0.5595	0										
662	6.1456	0.2788	0.6041	0										
750	6.5312	0.2919	0.6257	0	0.3	0.64	0.64	0.65	0.68	0.73	0.80	0.89	1.02	1.17
1000	7.5280	0.3245	0.6769	0										
1173	8.1472	0.3437	0.7067	0	0.5	0.69	0.69	0.69	0.71	0.75	0.81	0.89	1.00	1.13
1250	8.4384	0.3523	0.7198	0										
1333	8.7504	0.3617	0.7336	0	0.7	0.72	0.72	0.72	0.73	0.77	0.82	0.89	0.99	1.11
1460	9.1472	0.3731	0.7511	0										
1765	10.091	0.3997	0.7897	0	1.0	0.75	0.75	0.75	0.76	0.78	0.83	0.89	0.98	1.08
2004	10.818	0.4188	0.8173	C										
2250	11.397	0.4357	0.8414	C	1.5	0.78	0.78	0.78	0.79	0.81	0.84	0.89	0.96	1.05
2500	12.173	0.4536	0.8667	1	•••									
					2.0	0.80	0.80	0.81	0.82	0.82	0.85	0.89	0.95	1.02
					2.5	0.82	0.82	0.83	0.83	0.84	0.86	0.89	0.94	1.0

Health Physics Vol. 55, No. 1 (July), pp. 15-29, 1988 Printed in the U.S.A. 0017-9078/88 \$3.00 + .00 © 1988 Health Physics Society Pergamon Press pic

Paper

CALIBRATION FACTORS FOR Ge DETECTORS USED FOR FIELD SPECTROMETRY

Irene K. Helfer* and Kevin M. Miller** Environmental Measurements Laboratory, U.S. Department of Energy, New York, NY 10014-3621

(Received 26 June 1987; accepted 17 February 1988)

Abstract—Calibration factors to convert a measured full-absorption peak count rate to activity in the soil and dose rate in air are given for Ge detectors that are used for field measurements of radionuclides. The appropriate factors for a given detector are derived using three primary parameters: the manufacturer's quoted efficiency at 1332 keV relative to a 7.6 cm (3 in.) long by 7.6 cm (3 in.) diameter NaI(TI) detector, the detector's orientation in the field (up or down) and the Ge crystal length/diameter ratio. The accuracy of the results obtained by using this simplified calibration technique is estimated to be 10-15%.

INTRODUCTION

As a RESULT of the Three Mile Island incident in 1979, a need emerged for portable instrumentation that could be used for making rapid measurements in the field during emergency situations. Portable Ge detectors became available shortly thereafter and since have demonstrated their usefulness in the area of field spectrometry. These portable detectors have the same spectrometric capabilities as the standard large dewar types but are light weight and feature increased capabilities such as ruggedness, compactness and an all-attitude operation. With the advent of battery-powered portable microprocessor-based multichannel analyzers, a complete γ spectrometer system now can be hand carried by one person for use in remote areas.

Most γ spectrometrists are familiar with the laboratory calibration procedures used for fixed geometry sample analyses. However, for field work, they only may perform qualitative measurements by simply identifying the γ emitters present at a site. Semi-quantitative *in-situ* results are sometimes inferred by comparing peak count rates and obtaining a relative measure of contamination. Ideally, one would like to convert the count rate to some meaningful quantity such as dose rate or radionuclide concentration.

A Ge- γ spectrometer, whether portable or the standard large dewar-based type, has the capability to measure uncollided flux from various photon-emitting radionuclides in the environment. The Environmental Measurements Laboratory (EML) pioneered in studies involving

 Present Address: Law Associates, 2 University Plaza Suite 201, Hackensack, NJ 07601.

the application of *in-situ* γ spectrometry (Beck et al. 1972) to many types of radiation studies. These applications include the measurement of residual 137Cs levels in soils (Miller and Helfer 1985), an analysis of power plant reactor plumes (Gogolak 1984) and surveys of indoor exposure rates (Miller and Beck 1984). In-situ spectrometry proved to be extremely valuable in making rapid measurements of fission products in the environment during the Three Mile Island incident (Miller et al. 1979) and most recently was used with the same success by European laboratories during the Chernobyl crisis (Gogolak et al. 1986). In all of these applications, the measured photon flux can be converted into the inventory (activity per unit area) or concentration of a particular radionuclide in the soil and/or the exposure rate in the air based on a knowledge of the source distribution.

Since the adoption of Ge detectors for field spectrometry in 1971, EML has calibrated eight coaxial Ge detectors of various types and efficiencies. These were commercial units, purchased from two different manufacturers, that included Ge(Li) and high purity P-type and N-type Ge, ranging in efficiency from 3%-45% (at 1332 keV relative to a 7.6 cm [3 in.] long × 7.6 cm [3 in.] diameter Nal[TI] detector). The calibration data for these detectors have been reevaluated and correlated to the detector crystal dimensions and manufacturer's quoted efficiency. Based on this information, we present here analytical functions and tabular data that can be used to provide reasonably accurate field calibration factors for others to apply to their own Ge detectors.

FIELD SETUP PROCEDURES

In order to obtain accurate measurements of radionuclides in the soil, the Ge detector should be placed on

^{**} To whom correspondence should be addressed.



IAEA-TECDOC-1092

• 1999

IAEA-TECDOC-1092 Generic procedures for monitoring in a nuclear or radiological emergency PROCEDURE D.1.a, p 132



Generic procedures for monitoring in a nuclear or radiological emergency



INTERNATIONAL ATOMIC ENERGY AGENCY



June 1999

History

• *2013*

ISO 18589-7 Measurement of radioactivity in the environment — Soil — Part 7: In situ measurement of gamma-emitting radionuclides



INTERNATIONAL STANDARD

ISO 18589-7

First edition 2013-10-01

Measurement of radioactivity in the environment — Soil —

Part 7: In situ measurement of gammaemitting radionuclides

Mesurage de la radioactivité dans l'environnement — Sol — Partie 7: Mesurage in situ des radionucléides émetteurs gamma



Reference number ISO 18589-7:2013(E)

Laboratory vs In-Situ Gamma Measurement

	Laboratory measurement	In-Situ measurement
Sample information	Well known	Limited and/or estimated
Geometries	Small objects, bottles, beakers, Marinelli beakers	Surfaces, buildings, drums, boxes, large areas
Efficiency Calibration	Etalon available	Etalon not available
Results uncertainties	≈ < 5 %	≈ > 20 %

Object vs Ground Measurement

	Object measurement	Ground measurement
Space	Finite	Semi-Infinite
Sample	Container, Box, Package, Spill	Large plane areas
Results	Absolute or mass units Bq, Bq/kg	Surface or mass units Bq/m², Bq/kg

Background Radiation

- The nuclides usually measured by gamma spectrometry are the primordial nuclides: ⁴⁰K, ²³⁵U, ²³⁸U and ²³²Th.
- A few common reactor <u>activation and fission products</u> that are often present in background (¹³⁷Cs).
- There are, of course, other naturally occurring nuclides, such as ¹⁴C, which are produced continuously by nuclear reactions between high-energy particles with oxygen and nitrogen in the earth's atmosphere. Of those, only ⁷Be is measurable by gamma spectrometry.
- The major '<u>fluorescence' X-rays from likely shielding materials</u> Pb, Sn, Cd and Cu.

Background Gamma Spectrum



In-Situ Radionuclides

- **Primordial**
 - 40K
 - Decay chains/series
 - Uranium Radium ²³⁸U ²¹⁴Pb, ²¹⁴Bi \bullet
 - Thorium ²³²Th ²²⁸Ac, ²¹²Pb, ²⁰⁸Tl •
 - Actinium ²³⁵U •
- Artificial Nuclear weapons and \bullet reactors
 - ¹³¹I, ¹³⁴Cs, ¹³⁷Cs





²¹²Bi (60 min.)

35.94% α decay

(3.0 min)

β⁻ decay

64.06% β⁻ decay

α decay ²⁰⁸Pb (stable)

²¹²Po (0.299 μsec.)

228 Th (1.9 yr.)

α decay

α decay

Rn (55 sec.)

⁴Ra (3.6 day)

Background Radiation – ²³⁸U

- ²³⁸U comprises 99.25% of natural uranium. That decays by alpha emission to ²³⁴Th which in turn decays to ^{234m}Pa and so on until stable ²⁰⁶Pb is reached.
- If we look at the half-lives of the various nuclides, they are all much less than the half-life of ²³⁸U. This means that, in a natural, undisturbed source of uranium, every daughter nuclide will be in secular equilibrium with the ²³⁸U. The activity of each daughter nuclide will be equal to the ²³⁸U activity.
- There are 14 radionuclides in the chain and so the total activity of such a source will be 14 times that of the parent, or of any individual nuclide.

Background Radiation – ²³⁸U



Background Radiation – ²³⁵U

- ²³⁵U comprises 0.72 % of natural uranium.
- The decay series involves 12 nuclides in 11 decay stages and the emission of 7 alpha particles
- Within this series, only ²³⁵U itself can readily be measured.
- In the gamma spectra of NORM is the mutual interference between ²³⁵U (185.72 keV) and ²²⁶Ra (186.21 keV).
- Assume that the entire 186 keV peak is due to ²²⁶Ra and correct the result.

Corrected ²²⁶Ra = 0,5709×Apparent ²²⁶Ra Estimated ²³⁵U = 0,02662×Apparent ²²⁶Ra

Background Radiation – ²³²Th

- Natural thorium is 100 % ²³²Th. Six alpha particles are emitted during ten decay stages. Four nuclides can be measured easily by gamma spectrometry: ²²⁸Ac, ²¹²Pb, ²¹²Bi and ²⁰⁸Tl.
- The decay of ²¹²Bi is branched only 35.94 % of decays produce ²⁰⁸TI by alpha decay.
- If a ²⁰⁸TI measurement is to be used to estimate the thorium activity, it must be divided by 0.3594 to correct for the branching.

Background Radiation – ²³²Th



Background Radiation – ⁴⁰k

- ⁴⁰K is very evident in background spectra. It is present as <u>0.17% of</u> <u>natural potassium</u> and is present in wood and building materials and even in the bodies of the gamma spectrometrists.
- The substantial presence of ⁴⁰K in the detector background and in many samples, with its long Compton continuum, severely restricts the limit of detection of the many nuclides emitting gamma rays at lower energies
- 1460.83 keV

Measurement Conditions

- No buildings, forest, objects (at least 20 m - 30 m radius.
- No people (at least 5 metres)
- Vertical position of detector (down)
- 1 m detector/surface distance
- Energy > 60 100 keV





Detectors

• HPGe

- LN₂ detector + analyser + computer
- Portable electrically cooled
- Scintillation NaI:Tl or LaBr₃:Ce
 - Limited resolution

- Orientation of the detector
 - Down (preferable)
 - Up (laboratory configuration with Dewar)







Efficiency Calibration - 3 approaches

- Empirical approach based on the calibration standard measurements
 - Calibration based on using calibration standards (phantoms)
- Numerical approach (modelling) based on the numerical calculations
 - Possible to calculate any shape
 - Mathematical simulations (MCNP, Geant4, ISOCS SW)
- Combination of modelling and empirical approach
 - For example, ISOTOPIC SW (Pont source, DOE-EML % method)

Empirical Approach

- Spectra evaluation based on the Peak Net Area counts
- Calibration standard geometry = Sample geometry

$$\varepsilon(E) = \frac{N}{S \times t \times \gamma} \qquad A = \frac{N}{\varepsilon \times t}$$

$\underline{\varepsilon}(E)$ is efficiency,

- *N* net counts in full-energy peak area,
- *S* activity of certified calibration source
- γ emission probability of the gamma-ray with energy *E*
- t time of measurement
- *A* activity of sample

Modelling Approach

LaBr3(Ce) detector				
592 0	(-7.10.31):(-11	1 13 -32):(-10		
593 90 5.32	(-4 5 -32): 5	0 3 - 34		
595 90 - 7 596 902 - 2.7	(-3 4 -36):(- (-5 6 -56 -36):(-	1 3 -36) 6 6 36 -37):(-		
507.0	(-11 12 32 -37):	:(-12 13 32 -33	ע אך ו	
557 0	(-2 3 -39):(-3 ((-12 14 33 -39)		
598 902 -2.7 599 904 -0.0012759	-1 15 -40 (39: (1:-15:40)41	:2) imp:p 1 -999 imp:p 1		
588 903 -1	-41 imp:p 1	imara 0		
556 0	333	100.00		
c surface cards 1 pz 0				
2 pz -0.15				
4 pz -0.6	\$			-
5 pz -0.73 6 pz -1.36	\$			





Efficiency Calibration - Problems

- Calibration standard unavailable
- Distribution of radionuclides in the soil assumed and/or unavailable



- 3 cases, differ in depth distribution
 - Fresh Fallout
 - Aged Fallout
 - Homogeneous distribution



- The depth source distribution in the soil can be described by so-called α/ρ \bullet parameter or *B* parameter
- L [cm] relaxation length (length, where concentration decreases *e*-times) •
- α [cm⁻¹] 1 / relaxation length
- ρ [g cm⁻³] density

• α/ρ [cm² g⁻¹] α/ρ parameter • ß [g cm⁻²] relaxation mass per unit area, $\alpha/\rho = 1/\beta$ Uniform source distribution (NORM) $\alpha/\rho = 0$ or $\beta = \infty$ Surface source distribution in the soil (fresh fallout) $\alpha/\rho = \infty$ or $\beta = 0$ Exponential source distribution in the soil (aged fallout) $0 < \frac{\alpha}{\rho} < \infty$ or $0 < \beta < \infty$

Distribution	Uniform	Surface	Exponential
Result unit	Bq kg⁻¹	Bq m ⁻²	Bq m ⁻²
α/ρ	0	∞	$0 < \alpha/\rho < \infty$
ß	∞	0	0 < ß < ∞
Nuclides	NORM K-40 and decay chains	Fresh fallout Fission products	Aged fallout

 $\alpha/\rho = 0.206 \,[\text{cm}^2 \cdot \text{g}^{-1}] \approx \text{relaxation length of 3 cm, soil density 1,65 gcc - very frequently used}$ value for aged fallout

Activity distribution estimation

- Sampling depth profile
- Multiple line line to line method

Calibration Approach

BASED ON THE POINT SOURCES MEASUREMENTS AND MATHEMATICAL MODEL OF THE SAMPLE.

- Beck, H. L., DeCampo, J., and Gogolak, C. In Situ Ge(Li) And Nai(Tl) Gamma-Ray Spectrometry.. United States: N. p., **1972**. Web. doi:10.2172/4599415.
- Helfer I. K., Miller K. M. Calibration factors for Ge detectors used for field spectrometry. Health Phys. **1988** Jul;55(1):15-29. doi: 10.1097/00004032-198807000-00002. PMID: 3391774.
- IAEA-TECDOC-1092

Generic procedures for monitoring in a nuclear or radiological emergency

 ISO 18589-7 Measurement of radioactivity in the environment — Soil — Part 7: In situ measurement of gamma-emitting radionuclides

Efficiency Calibration Approach

• Beck, Helfer, Miller ...and older publications.

$$\varepsilon(E) = \frac{N_f}{A} = \frac{N_0}{\Phi} \times \frac{N_f}{N_0} \times \frac{\Phi}{A}$$



angular correction factor geometrical factor (incident flux arriving to the detector per unit inventory

response factor (intrinsic efficiency of the detector)

of the radionuclide in the soil)

Efficiency Calibration Approach

• ISO 18589-7

$$\boldsymbol{\varepsilon}(\boldsymbol{E}) = \frac{N_f}{A} = \boldsymbol{n_0} \times \mathbf{W} \times \mathbf{G}$$

- n_0 response factor (intrinsic efficiency of the detector)
- W angular correction factor
- **G** geometrical factor (incident flux arriving to the detector per unit inventory of the radionuclide in the soil)

Nomenclature Comparison

Older literature
$$\varepsilon(E) = \frac{N_f}{A} = n_0 \times W \times G$$

ISO $\varepsilon(E) = \frac{N_f}{A} = \frac{N_0}{\Phi} \times \frac{N_f}{N_0} \times \frac{\Phi}{A}$

RESPONSE FACTOR

ANGULAR CORRECTION FACTOR

GEOMETRICAL FACTOR

Peak count rate due to a unit primary photon flux density of Energy E incident on the detector along the detector axis (normal to the detector face) Based on the PS measurement

$$\frac{N_0}{\Phi} = \frac{\frac{P_i}{T}}{\frac{Y \times A}{4\pi r^2}} = \frac{4\pi r^2 \times P_i}{Y \times A \times T}$$

r [*cm*] is the source to detector distance

 P_i [-] is net count in the full energy peak for energy E

Y[-] is gamma yield per decay for photons of energy E

A [Bq] is activity of point source

T [s] is the measuring time

Based on Helfer & Miller relationships

 $a' = 2.689 + 0.4996 \times ln(\varepsilon_{rel}) + 0.0969 \times ln^2(\varepsilon_{rel})$

 $b' = -1.315 + 0.02044 \times \varepsilon_{rel} - 0.00012 \times \varepsilon_{rel}^2$

Correct formula

$\phi = \frac{Y \times A}{4\pi r^2}$	$e^{-\mu_a x} e^{-\mu_h h}$
Negligible	
$e^{-\mu_a x} e^{-\mu_h h} \approx$	× 1

 $\mu_{a} [cm^{-1}] \\ x [cm] \\ \mu_{h} [cm^{-1}] \\ h [cm]$

is linear attenuation coefficient for gammas in airis detector endcap to source crystal holderis linear attenuation coefficient for gam. in source holderis source holder thickness

Linear attenuation coefficient in air, r = 100 cm, air density 0.00129 gcc

= 100 keV	μ _a = 2.03 E-2 cm ⁻¹	$e^{-\mu_{a}x} = 0.980^{*}$
= 500 keV	μ _a = 1.12 E-2 cm ⁻¹	$e^{-\mu a^{\chi}} = 0.989^*$
= 1 MeV	μ _a = 8.15 E-3 cm ⁻¹	$e^{-\mu_a x} = 0.992^*$
= 5 MeV	μ _a = 3.55 E-3 cm ⁻¹	$e^{-\mu_a x} = 0.996^*$
= 10 MeV	μ _a = 2.67 E-3 cm ⁻¹	$e^{-\mu_a \chi} = 0.997^*$

*Ref.: https://physics.nist.gov/cgi-bin/Xcom/xcom2

- = distance from the source to the crystal effective centre [cm]:
 - for E > 1 MeV gamma rays crystal effective centre is approximately at the geometric centre of the crystal
 - ii. for E < 0.1 MeV gamma rays effective centre is approximately at the crystal face
 - iii. for energy range between those two values an estimation of average penetration has to be made based on the absorption coefficient of the crystal

$$r = \frac{1}{\mu} \cdot \frac{1 - e^{-\mu d} (\mu d + 1)}{1 - e^{-\mu d}} + d_{\circ} + x$$

- μ = attenuation coefficient in Ge detector at energy E [cm⁻¹]
- d = Ge crystal thickness [cm]

r

d_o = cap-to-crystal distance [cm].

Angular correction factor $\frac{N_f}{N_0}$ $\frac{N_f}{N_0} = \frac{\int R(\theta)\Phi(\theta)d\theta}{\int \Phi(\theta)d\theta}$

- It depends on the detector and source distribution in the soil
- Angular correction factor can be calculated from the measurement according to the equation
- ISO 18589-7: Ba-133 + Eu-152 or mixed gamma point sources





Angular correction factor $\frac{N_f}{N_0}$

 $\frac{N_f}{N_0} = \frac{\int R(\theta) \Phi(\theta) d\theta}{\int \Phi(\theta) d\theta} \qquad \qquad R(\theta) = \frac{\frac{P_\theta}{t_\theta}}{\frac{P_0}{t_0}}$

 $R(\theta)$ peak count rate for gamma rays of energy E at angle θ relative to count rate at angle $\theta = 0$

 $\Phi(heta)$ gamma ray flux at energy E at angle heta

 $\frac{P_{\theta}}{t_{\theta}}$

 $\frac{P_0}{t_0}$

- peak count rate for gamma rays of energy E at angle θ
- peak count rate for gamma rays of energy E at angle $\theta = 0$

Angular correction factor $\frac{N_f}{N_0}$

Dependent on crystal length/diameter ratio



Geometrical factor $\frac{\Phi}{A}$

- parameter Φ/A is incident flux arriving at the detector per unit activity of the source
- not dependent on detector, but it is function of soil density, soil composition, air attenuation and the <u>source distribution in</u> the soil.

$$\alpha/\rho = 0$$

$$\frac{\Phi}{S_{\gamma}/\rho} = \frac{1}{2} \frac{\rho_{S}}{\mu_{S}} \frac{\mu_{a}}{\rho_{a}} \rho_{a}h \left(\frac{e^{-(\mu_{a}/\rho_{a})\rho_{a}h}}{(\mu_{a}/\rho_{a})\rho_{a}h} - E_{1}\left(\frac{\mu_{a}}{\rho_{a}}\rho_{a}h\right) \right)$$
exponential
$$\frac{\Phi}{S_{0}} = \frac{1}{2}\left(E_{1}\left(\frac{\mu_{a}}{\rho_{a}}\rho_{a}h\right) - e^{\frac{\alpha}{\rho_{s}}\frac{\rho_{s}}{\mu_{s}}\frac{\mu_{a}}{\rho_{a}}\rho_{a}h} E_{1}\left(\left(1 + \frac{\alpha}{\rho_{s}}\frac{\rho_{s}}{\mu_{s}}\right)\frac{\mu_{a}}{\rho_{a}}\rho_{a}h\right) \right)$$

$$\alpha/\rho = \infty$$

$$\frac{\Phi}{S_{0}} = \frac{1}{2} E_{1}\left(\frac{\mu_{a}}{\rho_{a}}\rho_{a}h\right)$$

- 3 cases, differ in depth distribution
 - Fresh Fallout
 - Aged Fallout
 - Homogeneous distribution



Vertical		L cm							
distributi	on	1 cm	2 cm	3 cm	5 cm				
	1 cm	0.632	0.393	0.283	0.181				
	2 cm	0.233	0.239	0.204	0.149				
	3 cm	0.085	0.145	0.145	0.121				
	4 cm	0.032	0.088	0.104	0.100				
Depth	5 cm	0.011	0.053	0.075	0.081				
	10 cm	0.007	0.082	0.153	0.233				
	15 cm	x	X	0.036	0.085				
	20 cm	X	X	X	0.032				
	25 cm	X	X	X	0.011				



The effect of the detector position and soil density ¹³⁷Cs, L=3cm

Soil density: 1.05 gcc									
	95 cm	100 cm	105 cm						
Efficiency	1.34E-04	1.33E-04	1.32E-04						
a, Bq/m²	1771	1777	1786						
Soil density: 1.6 gcc									
a, Bq/m²		2317							

The effect of the soil density and relaxation length for ¹³⁷Cs, L=3cm

L, cm	1,05 kg/dm ³	1,6 kg/dm ³
1	1005	552
2	1276	1571
3	1777	2317
5	2302	3128

Real Depth Distribution

Vertical distribution of Cs-137 (%), at the site of in-situ measurement



ISOCS

- Efficiency calibration SW for In Situ Object Counting System
- Detector is already characterized by 5 step process
 - Traceable validation sources
 - Characterization measurements
 - MCNP Modelling
 - Detector characterization grid generation
 - Verification of characterization



Homogenous distribution - ISOCS



Exponential Distribution - ISOCS



Homogenous distribution-ISOCS



ISOTOPIC

- SW for measuring wide-area contamination of soils and surfaces
- Isotopic Supervisor (for measurement definition)
- Isotopic Operator (for analysis and reporting)
- Geometry Composer (a tool for measurement geometry definition)
- Two ways of efficiency calibration
 - DOE/EML Helfer and Miller estimation
 - Point Source measurement



ISOTOPIC

🔛 Soil Gamma Analysis Parameter Setup	- 🗆 ×	Soil Gar	mma Analysis Parameter Setup	- 🗆 X		
File Alpha/Rho Print Tables		File Alpha	a/Rho Print Tables			
Soil Setup File:	OK	Soil Setup	p File:	OK		
Field ID: ASDF	Cancel	Fie	Id ID: ASDF	Cancel		
			· · · · · · · · · · · · · · · · · · ·			
Detector Acquisition A	Analysis Heport		Analysis Analysis	- Heport		
001 uDetective	•					
Efficiency Calculation		Bea	al Time Preset: 3600			
DOE - EML Efficiency	_					
Detector Eff. 14.59 % Detector Height 100	cm	Livi	e Time Preset: U			
Crystal Len.: 36.1 (mm) Orientation: Down	•					
Crystal Dia.: 50.2 (mm) AspectRatio: 0.7						
C Line File: CALLAND REAMOND Schwarth E DANIA 2021	01 12 CALIERATION NO.		Soil Density: 1.6 g/cm^3			
S Geer IIIC. TC. 10581518 MAND 1085Klop1ME HANIA 12021	OT_T3_CALIBRATION_NO		Air Density: 0.0012 g/cm^3		DENSITIE	
Energy/FWHM Calibration						
Use File: C:\Users\BRANO\Desktop\MERANIA\2020	0_10_01_CALIBRATION_ST					
Energy: 0.531 + 3.6489e-1 * c + 1.2995e-7 * c**2						
FWHM: 2.915 + 7.6925e-4 * c + -2.7942e-8 * c**2	🔄 Alpha/Rho Value			Soil Gamma Ana	lysis Parameter Setup	- 🗆 X
		Nuclide:		File Alpha/Rho	rint Tables	
		100 00		Soil Setup File:		
	Units	ource Distribution	Cancel AL DILLA	Field ID: A	3DF	Cancel
	C Default	🔿 Uniform (Alpha/Rho = 0)		Detector	Acquisition Analysis	Report
	C g 🔤	O Plane (Alpha/Rho = Infinity)	RHO —			
	⊙ m^2	Exponential (Alpha/Rho >= .0625)		Alpha /Rho File:		
		Alpha/Bho = 1	FILE		verride Value 1 🔽 🔽 Optimize Alpha/Rho	
		en Conservationes Inc.		Library File:	C:\Users\BBAND\Deskton\MEBANIA\2020_12_21_IN_SI	
	- Concentration Batia Limite		SOURCE	Libidiy File. J		
				Analysis Engine:	IsoEnv32.Exe Directed Fit	
	Energy 1 (keV)	Energy 2 (keV) Lower Limit Upper Li			70 % Decay Date:	
	1173.23	1332.51 0.5 10			dd/MM/yyyy	
CALIBRATION				Match Width:	2 × FWHM Analyze Chan 10 to 8000	
	Energy 1: 1173.23	keV Lower Limit: 0,5		Park Ostaffi	*	- I
	Energy 2: 1332.51	keV Upper Limit: 10		reak cuton:	UU ~ Background Type: 5 points _	
	Add	Update Delete				

Pros & Cons

PROS

FAST MEASUREMENT NO SAMPLE

FAST RESULTS

CONS

HIGH UNCERTAINTIES

DETECTOR CONTMINATION POSSIBILITY

ACTIVITY DISTRIBUTION DETERMINATION

CASE STUDY – HOMOGENOUS DISTRIBUTION Example Ac-228 Manual Calibration

Efficiency Calibration Approach

Beck, Helfer, Miller \bullet

$$\varepsilon(E) = \frac{N_f}{A} = \frac{N_0}{\Phi} \times \frac{N_f}{N_0} \times \frac{\Phi}{A}$$

- $\frac{N_0}{\Phi}$ response factor (intrinsic efficiency of the detector)
- $\frac{\bar{N_f}}{N_0}$ angular correction factor
- $\frac{\Phi}{A}$ geometrical factor (incident flux arriving to the detector per unit inventory of the radionuclide in the soil)





 Peak counting rate due to a unit primary photon flux density of Energy E incident on the detector along the detector axis (normal to the detector face)

 $I \mid S \mid$

- Point Sources at distance of 100 cm
- Whole energy region of interest

$$\frac{N_{0}}{\Phi} = \frac{\frac{P_{i}}{T}}{\frac{Y \times A_{CALIB}}{4\pi r^{2}}} = \frac{4\pi r^{2} \times P_{i}}{Y \times A \times T} \quad \begin{array}{c} r \left[m\right] \\ P_{i}\left[-\right] \\ Y\left[-\right] \\ A_{CALIB}\left[Bq\right] \\ T_{i}\left[-\right] \end{array}$$

is the source to detector distance
is net count in the full energy peak for energy E
is gamma yield per decay for photons of energy E
is activity of point source
is the measuring time

- Any SW
- Peak net Area



 \bigcirc

 $\frac{\Psi_0}{\Phi}$

 $\varepsilon(E)$

 $\varepsilon(E) = \frac{N_0}{\Phi} \times \frac{N_f}{N_0} \times \frac{\Phi}{A}$

Energy [keV]	121	244	344	661	778	898	964	1085	1112	1173	1332	1408	1836
Nuclide	152-Eu	152-Eu	152-Eu	137-Cs	152-Eu	Y-88	152-Eu	152-Eu	152-Eu	60-Co	60-Co	152-Eu `	Y-88
A [kBq]	450	450	450	591	450	47	450	450	450	417	417	450	47
T [s]	3600	3600	3600	3600	3600	3600	3600	3600	3600	3600	3600	3600	3600
Y[-]	0.284	0.076	0.266	0.850	0.130	0.937	0.145	0.101	0.134	0.999	1.000	0.209	0.993
N [-]	84983	16758	46801	113793	12064	8113	11743	7265	9918	57725	53349	12985	5019
N0 / Φ [cm ²]	23.665	17.560	13.925	8.112	7.359	6.569	6.407	5.674	5.851	5.274	4.868	4.927	3.833





- It depends on the detector and source distribution in the soil.
- Angular correction factor can be calculated from the measurement according to the equation.
- This factor expresses weighted angular response compared to the normal response

Angular correction factor $\frac{Nf}{Nc}$



• ISO

$$W = \sum_{m=1}^{M} k_m \cdot \left(\frac{\Delta \Phi_m}{\Phi}\right)$$



Portion of flux density of unscattered photons of energy *E* resulting from polar angle segment m for distribution model *V* at the detector location

$$k_m = \frac{\eta_m}{\eta_0}$$

η_m	Cross section of the detector for photons from the polar seg-	m^2
	ment, <i>m</i>	

 η_0 Intrinsic efficiency

 ${\rm m}^2$

Angular correction factor $\frac{N_f}{N_f}$



HOMOGENOUS DISTRIBUTION

$$\left(\frac{\Delta \Phi_m}{\Phi}\right)_{E,V} = \frac{E_2(\mu_{Air} \cdot d / \cos \vartheta_{int}) \cdot \cos \vartheta_{int} - E_2(\mu_{Air} \cdot d / \cos \vartheta_{ext}) \cdot \cos \vartheta_{ext}}{E_2(\mu_{Air} \cdot d)}$$

$$E_1$$
 1. order exponential integral function $E_1(\alpha) = \int_{1}^{\infty} \frac{e^{-\alpha x}}{x} dx$

*E*₂ 2. order exponential integral function
$$E_2(\alpha) = \int_{1}^{\infty} \frac{e^{-\alpha x}}{x^2} dx$$

Angular correction factor $\frac{N}{N}$

Deposits on the ground surface:

$$\left(\frac{\Delta \Phi_m}{\Phi}\right)_{E,V} = \frac{E_1(\mu_{Air} \cdot d / \cos \vartheta_{int}) - E_1(\mu_{Air} \cdot d / \cos \vartheta_{ext})}{E_1(\mu_{Air} \cdot d)}$$

Exponential distribution in soil:

$$\left(\frac{\Delta \Phi_{m}}{\Phi}\right)_{E,V} = \frac{E_{1}(\mu_{Air} \cdot d / \cos \vartheta_{int}) - \exp\left(\mu_{Air} \cdot d \cdot \frac{1/\beta}{\mu_{S}/\rho_{S}}\right) \cdot E_{1}\left(\mu_{Air} \cdot d \cdot \left(\frac{1/\beta}{\mu_{S}/\rho_{S}} + d / \cos \vartheta_{int}\right)\right)}{E_{1}(\mu_{Air} \cdot d) - \exp\left(\mu_{Air} \cdot d \cdot \frac{1/\beta}{\mu_{S}/\rho_{S}}\right) \cdot E_{1}\left(\mu_{Air} \cdot d \cdot \left(\frac{1/\beta}{\mu_{S}/\rho_{S}} + 1\right)\right)}$$
$$-\frac{E_{1}(\mu_{Air} \cdot d / \cos \vartheta_{ext}) - \exp\left(\mu_{Air} \cdot d \cdot \frac{1/\beta}{\mu_{S}/\rho_{S}}\right) \cdot E_{1}\left(\mu_{Air} \cdot d \cdot \left(\frac{1/\beta}{\mu_{S}/\rho_{S}} + d / \cos \vartheta_{ext}\right)\right)}{E_{1}(\mu_{Air} \cdot d) - \exp\left(\mu_{Air} \cdot d \cdot \frac{1/\beta}{\mu_{S}/\rho_{S}}\right) \cdot E_{1}\left(\mu_{Air} \cdot d \cdot \left(\frac{1/\beta}{\mu_{S}/\rho_{S}} + d / \cos \vartheta_{ext}\right)\right)}\right)$$

 $\varepsilon(E)$

Uniform distribution in soil:

$$\left(\frac{\Delta \Phi_m}{\Phi}\right)_{E,V} = \frac{E_2(\mu_{Air} \cdot d / \cos \vartheta_{int}) \cdot \cos \vartheta_{int} - E_2(\mu_{Air} \cdot d / \cos \vartheta_{ext}) \cdot \cos \vartheta_{ext}}{E_2(\mu_{Air} \cdot d)}$$

Angular correction factor

- not so simple calculation.....
- Hellfer and Miller tables, Isotopic SW manual, etc.....
 - Angular Correction Factor, (Nf /N0), Downward-Facing Detector, Plane Source Distribution (α/ρ =0).

E	nergy	L/D									
[]	NeV]	0.5	0.6	0.7	0.8	0.9	1.0	1.1	1.2	1.3	
	0.3	0.81	0.82	0.83	0.86	0.91	0.99	1.08	1.18	1.31	
	0.5	0.84	0.85	0.85	0.88	0.93	0.99	1.06	1.14	1.25	
	0.7	0.86	0.86	0.87	0.91	0.93	0.98	1.05	1.12	1.21	
	1.0	0.88	0.88	0.89	0.91	0.94	0.98	1.03	1.10	1.18	
	1.5	0.91	0.91	0.91	0.92	0.94	0.97	1.02	1.07	1.13	
	2.0	0.92	0.92	0.93	0.93	0.94	0.96	1.00	1.05	1.10	
	2.5	0.94	0.94	0.94	0.94	0.95	0.96	0.99	1.03	1.07	



 $\varepsilon(E)$

Geometrical factor

- $\boldsymbol{\varepsilon}(\boldsymbol{E}) = \frac{N_0}{\boldsymbol{\Phi}} \times \frac{N_f}{N_0} \times \frac{\boldsymbol{\Phi}}{\boldsymbol{A}}$
- The last parameter Φ/A is incident flux at that energy arriving at the detector per unit inventory or concentration of the nuclide in the soil. It is not dependent on detector, but it is function of soil density, soil composition, air attenuation and

the source distribution in the soil.

$$\alpha/\rho = 0$$

$$\frac{\Phi}{S_{\gamma}/\rho} = \frac{1}{2} \frac{\rho_{S}}{\mu_{S}} \frac{\mu_{a}}{\rho_{a}} \rho_{a}h \left(\frac{e^{-(\mu_{a}/\rho_{a})\rho_{a}h}}{(\mu_{a}/\rho_{a})\rho_{a}h} - E_{1}\left(\frac{\mu_{a}}{\rho_{a}}\rho_{a}h\right) \right)$$
exponential
$$\frac{\Phi}{S_{0}} = \frac{1}{2} \left(E_{1}\left(\frac{\mu_{a}}{\rho_{a}}\rho_{a}h\right) - e^{\frac{\alpha}{\rho_{s}}\frac{\rho_{s}}{\mu_{s}}\frac{\mu_{a}}{\rho_{a}}\rho_{a}h} E_{1}\left(\left(1 + \frac{\alpha}{\rho_{s}}\frac{\rho_{s}}{\mu_{s}}\right)\frac{\mu_{a}}{\rho_{a}}\rho_{a}h\right) \right)$$

$$\alpha/\rho = \infty$$

$$\frac{\Phi}{S_{0}} = \frac{1}{2} E_{1}\left(\frac{\mu_{a}}{\rho_{a}}\rho_{a}h\right)$$

Geometrical factor

 Beck established the following table for unscattered flux one meter above the ground for various energies and α/ρ values.

$[(cm^{-2} s^{-1}) / Bq q^{-1})]$ for $\alpha/\rho = 0$ (uniform profile, natural emitters)								
$[(cm^{-2} s^{-1}) / Bq cm^{-1})]$ for $\alpha/\rho > 0$ (exponential profile, fallout)								
Energy	Ó	α/ρ			·	,		
[MeV]		0	0.06	0.206	0.312	0.625	6.25	∞
(0.050	1.4403	0.0816	0.2245	0.3049	0.4748	1.147	1.58
	0.100	2.1140	0.1458	0.3627	0.4708	0.6786	1.359	1.71
	0.150	3.3264	0.1702	0.4103	0.5261	0.7438	1.427	1.78
	0.200	3.9056	0.1843	0.4550	0.5770	0.8020	1.483	1.80
	0.250	4.0640	0.2008	0.4697	0.5910	0.8185	1.506	1.86
	0.364	4.7184	0.2268	0.5158	0.6429	0.8775	1.578	1.93
	0.500	5.3904	0.2519	0.5595	0.6918	0.9334	1.650	2.00
	0.662	6.1456	0.2788	0.6041	0.7412	0.9889	1.719	2.05
	0.750	6.5312	0.2919	0.6257	0.7649	1.0150	1.752	2.08
	1.000	7.5280	0.3245	0.6769	0.8209	1.0770	1.830	2.15
	1.173	8.1472	0.3437	0.7067	0.8531	1.1130	1.874	2.19
	1.250	8.4384	0.3523	0.7198	0.8675	1.1290	1.895	2.21
	1.333	8.7504	0.3617	0.7336	0.8826	1.1450	1.914	2.22
	1.460	9.1472	0.3731	0.7511	0.9011	1.1660	1.941	2.25
	1.765	10.0910	0.3997	0.7897	0.9428	1.2110	1.997	2.29
	2.004	10.8180	0.4188	0.8173	0.9725	1.2430	2.036	2.33
	2.250	11.3970	0.4357	0.8410	0.9982	1.2710	2.071	2.36
	2 500	12,1700	0 4536	0 8667	1 0250	1 3000	2 105	2 39

 $\mathcal{E}(\underline{E})$

Peak Net Area – ²²⁸Ac



Efficiency Calibration Approach – ²²⁸Ac

Energy [keV]	338.3	911.2	964.8	969.0
N ₀ / Φ [cm²]	14.10	6.62	6.35	6.33
N_f/N_0 [-]	0.99	0.98	0.98	0.98
Φ / A [cm ⁻² g]	4.75	7.07	7.29	7.31
N ₀ /A [g]	66.29	45.86	45.33	45.30



Results – ²²⁸AC $A = \frac{N}{T * Y * \varepsilon}$

A [Bq kg-1]	20.09	22.07	19.03	23.35
T [s]	7200	7200	7200	7200
Peak Net Area	1093	1909	310	1211
ε [g]	66.29	45.86	45.33	45.30
Y	0.114	0.262	0.049	0.159
Energy [keV]	338.3	911.2	964.8	969.0

CASE STUDY – HOMOGENOUS – RESULTS Example Ac-228

DETECTOR	METHOD	ACTIVITY
AEGIS	ISOCS	26.15 ± 0.89 Bq/kg
AEGIS	ISOTOPIC PS	23.92 ± 2.25 Bq/kg
AEGIS	ISOTOPIC %	20.87 ± 0.94 Bq/kg
AEGIS	MANUAL	21.14 ± 1.68 Bq/kg
LAB	LabSOCS	21.30 ± 0.40 Bq/kg

Thank you for the attention!



