Update of the BIPM comparison BIPM.RI(II)-K1.Sn-113 of activity measurements of the radionuclide ^{113}Sn to include the 2017 result of the LNE-LNHB (France)

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Abstract Since 1975, 7 laboratories have submitted 13 samples of ^{113}Sn to the International Reference System (SIR) for activity comparison at the Bureau International des Poids et Mesures (BIPM), with comparison identifier BIPM.RI(II)-K1.Sn-113. Recently, the LNE-LNHB (France) participated in the comparison and the key comparison reference value (KCRV) has been updated to take account of the change of the ^{113}Sn half-life value used for the SIR. The degrees of equivalence between each equivalent activity measured in the SIR and the updated KCRV have been calculated and the results are given in the form of a table. A graphical presentation is also given.

1. Introduction

The SIR for activity measurements of γ-ray-emitting radionuclides was established in 1976. Each national metrology institute (NMI) may request a standard ampoule from the BIPM that is then filled with 3.6 g of the radioactive solution. For radioactive gases, a different standard ampoule is used. Each NMI completes a submission form that details the standardization method used to determine the absolute activity of the radionuclide and the full uncertainty budget for the evaluation. The ampoules are sent to the BIPM where they are compared with standard sources of ^{226}Ra using pressurized ionization chambers. Details of the SIR method, experimental set-up and the determination of the equivalent activity A_e, are all given in [1].

From its inception until 31 December 2021, the SIR has been used to measure 1033 ampoules to give 788 independent results for 72 different radionuclides. The SIR makes it possible for national laboratories to check the reliability of their activity measurements

at any time. This is achieved by the determination of the equivalent activity of the radionuclide and by comparison of the result with the key comparison reference value determined from the results of primary standardizations. These comparisons are described as BIPM continuous comparisons and the results form the basis of the BIPM key comparison database (KCDB) of the Comité International des Poids et Mesures Mutual Recognition Arrangement (CIPM MRA) [2]. The comparison described in this report is known as the BIPM.RI(II)-K1.Sn-113 key comparison. The results of earlier participations in this key comparison were published previously [3, 4].

2. Participants

Laboratory details are given in Table 1, with the earlier submissions being taken from [3, 4]. The dates of measurement in the SIR given in Table 1 are used in the KCDB and all references in this report.

Table 1: Details of the participants in the BIPM.RI(II)-K1.Sn-113.

NMI or laboratory	**Previous acronyms**	**Full name**	**Country**	**RMO**	**Date of SIR measurement** yyyy-mm-dd
BKFH	OMH, MKEH	Government Office of the Capital City Budapest	Hungary	EURAMET	1988-09-19
CIEMAT	-	Centro de Investigaciones Energéticas, Medioambientales y Tecnologicas	Spain	EURAMET	2011-12-07
CMI	UVVVR, CMI-IIR	Czech Metrological Institute	Czechia	EURAMET	1981-12-16
CNEA	-	Comision Nacional de Energia Atomica	Argentina	SIM	1994-02-28
LNE-LNHB	LMRI, LPRI, BNM-LNHB	Laboratoire National de métrologie et d´Essais -Laboratoire National Henri Becquerel	France	EURAMET	1992-02-21 2010-08-02 2017-09-08
NIST	NBS	National Institute of Standards and Technology	United States	SIM	1975-09-15 1980-10-24
PTB	-	Physikalisch-Technische Bundesanstalt	Germany	EURAMET	1989-04-12 2010-03-31

3. NMI standardization methods

Each NMI that submits ampoules to the SIR has measured the activity either by a primary standardization method or by using a secondary method, for example a calibrated ionization chamber. In the latter case, the traceability of the calibration needs to be

clearly identified to ensure that appropriate correlations are taken into account.

A brief description of the standardization methods used by the laboratories, the activities submitted, the relative standard uncertainties and the half-life used by the participants are given in Table 2. The uncertainty budget for the new submission is given in Appendix D attached to this report; previous uncertainty budgets are given in the earlier K1 reports [3, 4]. The list of acronyms used to summarize the methods is given in Appendix E.

The half-life used previously by the BIPM was 114.9(1) days [5] and this was changed to the more recent evaluation of 115.09(3) days [6]. All the results in Table 4 have been updated accordingly. The impurity corrections for the SIR measurements were also updated.

Table 2: Standardization methods of the participants for ^{113}Sn.

NMI or laboratory	**Method used and the acronym**	**Activity A_i/kBq**	**Relative standard uncertainty /10^{-2}**		**Reference date**	**Half-life /d**
			A	**B**	**yyyy-mm-dd**	
BKFH	Ge(Li) γ counting (UA-GL-GR-00-00-00)	10 350	0.2	1.0	1988-10-01 12:00 UT	115.08(3) [7]
CIEMAT	4π NaI(Tl) counting (4P-NA-GR-00-00-HE)	724.2	0.10	0.61	2011-10-20 10:00 UT	115.09(3) [6]
	4πec-x coincidence (4P-PP-CE-NA-XR-CO)	724.8	0.87	0.24		
CMI	4πec-x coincidence (4P-PC-CE-NA-XR-CO)	3922[g]	0.13	0.7	1981-11-24 11:00 UT	115.1
	4πec counting (4P-PC-CE-00-00-00)					
CNEA	HP-Ge spectrometry (UA-GH-GR-00-00-00)	914	0.28	0.76	1994-01-01 12:00 UT	115.09
LNE-LNHB	HP-Ge spectrometry (UA-GH-GR-00-00-00)[a]	1135.9[f]	0.03	0.36	1992-01-28 12:00 UT	115.08(3) [7]
		1138.4	0.03	0.36		
	4π NaI(Tl) counting (4P-NA-PH-00-00-HE)	1175.5	0.41	0.32	2010-07-01 12:00 UT	115.09(3) [6]
	Gamma spectrometry (UA-GH-GR-00-00-00)[b]	503.5	0.59	0.61	2017-10-09 12:00 UT	[6]
NIST	ionization chamber (4P-IC-GR-00-00-00)[c]	7891	2.9		1975-07-01 17:00 UT	- [8]
	ionization chamber (4P-IC-GR-00-00-00)[c]	4738	0.01	1.07	1980-10-15 19:00 UT	
PTB	ionization chamber (4P-IC-GR-00-00-00)[d]	21 620	0.2	1.0	1989-02-01 00:00 UT	

... Continuation of Table 2.

NMI or laboratory	**Method used and the acronym**	**Activity A_i/kBq**	**Relative standard uncertainty /10^{-2}**		**Reference date** yyyy-mm-dd	**Half-life /d**
			A	**B**		
	CIEMAT/NIST (4P-LS-MX-00-00-CN) TDCR (4P-LS-MX-00-00-TD)	1258.4[e]	0.15	1.25	2010-03-01 00:00 UT	115.09(3) [6]

[a] Calibrated in 1990 using about 80 energy points not including those of Sn-113 γ transitions
[b] Calibrated using LNE-LNHB primary standards on a large energy range (> 20 radionuclides)
[c] Calibrated for Sn-113 by NaI and Ge(Li) photopeak counting UA-NA-GR-00-00-00 and UA-GL-GR-00-00-00
[d] Calibrated in 1980 by efficiency calculation and Ge spectrometry UA-GL-GR-00-00-00
[e] The result is the mean of the different methods. The uncertainty of CIEMAT/NIST method has been attributed to the weighted mean
[f] Several samples submitted
[g] The result is the mean of the different methods.

Details regarding the solutions submitted are shown in Table 3, including any impurities, when present, as identified by the laboratories. When given, the standard uncertainties on the evaluations are shown.

Table 3: Details of each solution of ^{113}Sn submitted.

NMI or laboratory / SIR year	**Chemical composition**	**Solvent conc.** /(mol dm^{-3})	**Carrier conc.** /(µg g^{-1})	**Density** /(g cm^{-3})	**Relative activity of any impurity[a]**
BKFH 1988	Sn + In in HCl	6	Sn: 28 In: 17	-	^{114m}In: 0.056(5) % ^{124}Sb: 0.0010(3) %
CIEMAT 2011	$SnCl_4$ in HCl	6	$SnCl_4$: 10	1.09	-
CMI 1981	Hexachloro-stannate in HCl	6	-	-	^{114m}In: 0.90(5) % ^{125}Sb: 0.040(5) %
CNEA 1994	$SnCl_2$	-	Sn: 32.6	0.9969	-
LNE-LNHB 1992	$SnCl_4$ in HCl	2	$SnCl_4$: 740	1.035	^{125}Sb: 0.06(1) % ^{60}Co: 0.002(1) %
2010	$SnCl_4$ in HCl	6	Sn: 10	1.098	^{114m}In: 0.051(8) % ^{110m}Ag: 0.011(1) %
2017	Sn(IV) in HCl	6	Sn: 10	1.098	^{110m}Ag: < 0.012 % ^{114m}In: < 0.021 %
NIST 1975	$SnCl_4$ in HCl	4	-	1.064(2)	-
1980	Sn in HCl	4	Sn: 2	1.066(2)	^{114m}In: 3.6(7) % ^{125}Sb: 0.12(2) %

... Continuation of Table 3.

NMI or laboratory / SIR year	**Chemical composition**	**Solvent conc. /(mol dm^{-3})**	**Carrier conc. /(µg g^{-1})**	**Density /(g cm^{-3})**	**Relative activity of any impurity[a]**
PTB 1989 2010	$SnCl_2$ in HCl	3	$SnCl_2$: 50	1.05	-
	$SnCl_2.2H_2O$ in HCl	4	$SnCl_2.2H_2O$: 47.4	1.05	^{114m}In: 0.15(5) % ^{75}Se: 0.022(9) % ^{110m}Ag: 5.7(14)x10^{-3} %

[a] The ratio of the activity of the impurity to the activity of ^{113}Sn at the reference date

4. Results

All the submissions to the SIR since its inception in 1976 are maintained in a dedicated database [9]. The latest submission has added 1 ampoule for the activity measurements for ^{113}Sn giving rise to 13 ampoules in total.

The SIR equivalent activity, A_{ei}, for each ampoule received from each NMI, i, including both previous and new results, is given in Table 4. The relative standard uncertainties arising from the measurements in the SIR are also shown. This uncertainty is additional to that declared by the NMI ($u(A_i)$) for the activity measurement shown in Table 2. Although submitted activities are compared with a given source of ^{226}Ra, all the SIR results are normalized to the radium source number 5 [1]. Table 4 also shows the comparison results selected for the KCRV as explained in section 4.1.

The assumption is made that the daughter radionuclide ^{113}In (half-life of 99 min) was in equilibrium with the parent at the reference dates.

In principle, the chemical composition of the solutions could have an influence on the SIR measurements owing to the intense x-ray emission from ^{113}Sn. However, using the efficiency curve of the SIR [10], the contribution of the x-rays to the ionization current is estimated to be negligible. In consequence, the influence of the chemical composition on the SIR measurements can also probably be neglected in this case although a more detailed study could be performed.

Table 4: Results of SIR measurement of ^{113}Sn.

NMI or laboratory / SIR year	m_i **/g**	A_i **/kBq**	^{226}**Ra source**	A_{ei} **/kBq**	**Relative uncert. from SIR /10^{-4}**	u_{ci} **/kBq**	A_{ei} **for KCRV /kBq**
BKFH 1988	3.650 2	10 350	4	58 780	7	620	-

... Continuation of Table 4.

NMI or laboratory / SIR year	m_i **/g**	A_i **/kBq**	^{226}**Ra source**	A_{ei} **/kBq**	**Relative uncert. from SIR** $/10^{-4}$	$u_{c,i}$ **/kBq**	A_e **for KCRV /kBq**
CIEMAT 2011	3.621 3	724.2[a] 724.8	1	58 430[b]	15	310	58 470(540)[b2] -
CMI 1981	3.634 91	3922	3	58 970	8	420	58 970(420)
CNEA 1994	3.672 32	914	1	58 380	13	480	-
LNE-LNHB 1992	3.703 5	1135.9	2	58 269	11	220	-
	3.711 6	1138.4	2	58 321	10	220	-
2010	3.976 8(6)[c]	1175.5	1	57 130	15	310	-
2017	4.015 2[c]	503.5	1	58 790	16	510	-
NIST 1975	5.368 1[d]	7891	3	58 980[e]	9	1700	-
1980	3.865 63	4738	3	58 870	23[f]	640	-
PTB 1989	3.628 7	21 620	4	58 250	7	600	-
2010	3.821 33	1258.4	1	59 110	16	750	59 110(750)

[a] Activity values corresponding to the two methods mentioned in Table 2.

[b] SIR result evaluated using the weighted mean activity measurement result as computed by the CIEMAT: 724.4(37) kBq

[b2] Using the coincidence method result only

[c] Mass higher than the limit of 3.8 g for SIR ampoules

[d] Masses higher than the limit of 3.8 g for SIR ampoules; part of the solution (3.9096 g) was transfered to another ampoule at the BIPM, giving an SIR result of 58 790(1700) kBq.

[e] The half-life used for these SIR measurements is 114.9(1) d.

[f] Mainly due to the uncertainty of the impurity activities as given by the NIST

4.1. The key comparison reference value

In May 2013, the CCRI(II) decided to calculate the key comparison reference value (KCRV) by using the power-moderated weighted mean [11] rather than an unweighted mean, as had been the policy. This type of weighted mean is similar to a Mandel-Paule mean in that the NMIs' uncertainties may be increased until the reduced chi-squared value is one. In addition, it allows for a power α smaller than two in the weighting factor. As proposed in [11], α is taken as $2 - 3/N$ where N is the number of results selected for the KCRV. Therefore, all SIR key comparison results can be selected for the KCRV with the following provisions:

(a) only results for solutions standardized by primary techniques are accepted, with the exception of radioactive gas standards (for which results from transfer instrument measurements that are directly traceable to a primary measurement in the laboratory may be included);

(b) each NMI or other laboratory may only use one result (normally the most recent result or the mean if more than one ampoule is submitted);

(c) results more than 20 years old are included in the calculation of the KCRV but are not included in data shown in the KCDB or in the plots in this report, as they have expired;

(d) possible outliers can be identified on a mathematical basis and excluded from the KCRV using the normalized error test with a test value of 2.5 and using the modified uncertainties;

(e) results can also be excluded for technical reasons; and

(f) the CCRI(II) is always the final arbiter regarding excluding any data from the calculation of the KCRV.

The data set used for the evaluation of the KCRVs is known as the KCRV file and is a reduced data set from the SIR master-file. Although the KCRV may be modified when other NMIs participate, on the advice of the Key Comparison Working Group of the CCRI(II), such modifications are made only by the CCRI(II) during one of its biennial meetings, or by consensus through electronic means (e.g., email) as discussed at the CCRI(II) meeting in 2013.

Consequently, using the results updated for the half-life produces an updated KCRV for ^{113}Sn in 2017 of **58 840(310) kBq** with the power α =1.0 that has been calculated using the previously published results, selected as shown in Table 4, for the CMI (1981), PTB (2010), and the CIEMAT (2011) result. This can be compared with the previous KCRV value 58 810(310) kBq published in 2016 [4].

4.2. Degrees of equivalence

Every participant in a comparison is entitled to have one result included in the KCDB as long as the NMI is a signatory or designated institute listed in the CIPM MRA and the result is valid (i.e., not older than 20 years). No recent submission has been identified as a pilot study so the most recent result of each NMI is normally eligible for inclusion on the KCDB platform of the CIPM MRA [2]. An NMI may withdraw its result only if all other participants agree.

The degree of equivalence of a given measurement standard is the degree to which this standard is consistent with the KCRV [2]. The degree of equivalence is expressed quantitatively in terms of the deviation from the key comparison reference value and the expanded uncertainty of this deviation ($k = 2$). The degree of equivalence between any pair of national measurement standards is expressed in terms of their difference and the expanded uncertainty of this difference and is independent of the choice of key comparison reference value.

4.2.1. Comparison of a given NMI result with the KCRV

The degree of equivalence of the result of a particular NMI, i, with the key comparison reference value is expressed as the difference D_i between the values

$$D_i = A_{ei} - \text{KCRV} \tag{1}$$

and the expanded uncertainty ($k = 2$) of this difference, U_i, known as the equivalence uncertainty; hence

$$U_i = 2u(D_i) \tag{2}$$

When the result of the NMI i is included in the KCRV with a weight w_i, then

$$u^2(D_i) = (1 - 2w_i)u_i^2 + u^2(\text{KCRV}) \tag{3}$$

However, when the result of the NMI i is not included in the KCRV, then

$$u^2(D_i) = u_i^2 + u^2(\text{KCRV}) \tag{4}$$

4.2.2. Comparison between pairs of NMI results

The degree of equivalence between the results of any pair of NMIs, i and j, is expressed as the difference D_{ij} in the values

$$D_{ij} = D_i - D_j = A_{ei} - A_{ej} \tag{5}$$

and the expanded uncertainty ($k = 2$) of this difference, $U_{ij} = 2u(D_{ij})$, where

$$u^2(D_{ij}) = u_i^2 + u_j^2 - 2u(A_{ei}, A_{ej}) \tag{6}$$

where any obvious correlations between the NMIs (such as a traceable calibration, or correlations normally coming from the SIR or from the linking factor in the case of linked comparison) are subtracted using the covariance $u(A_{ei}, A_{ej})$ (see [12] for more detail). However, the CCRI decided in 2011 that these pair-wise degrees of equivalence no longer need to be published as long as the methodology is explained.

Table B1 shows the matrix of all the degrees of equivalence as they will appear in the KCDB. It should be noted that for consistency within the KCDB, a simplified level of nomenclature is used with A_{ei} replaced by x_i. The introductory text is that agreed for the comparison. The graph of the results in Table 5, corresponding to the degrees of equivalence with respect to the KCRV (identified as x_R in the KCDB), is shown in Figure C1. This graphical representation indicates in part the degree of equivalence between the NMIs but obviously does not take into account the correlations between the different NMIs. It should be noted that the final data in this paper, while correct at the time of publication, will become out-of-date as NMIs make new comparisons. The formal results under the CIPM MRA [2] are those available in the KCDB.

5. Conclusion

The BIPM continuous key comparison for ^{113}Sn, BIPM.RI(II)-K1.Sn-113, currently comprises 3 results. The results have been analyzed with respect to the updated KCRV, providing degrees of equivalence for 3 national metrology institutes. The degrees of equivalence have been approved by the CCRI(II) and are published in the BIPM key comparison database. Other results may be added when other NMIs contribute ^{113}Sn activity measurements to this comparison or take part in other linked comparisons.

6. References

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Appendix A. Introductory text for ^{113}Sn degrees of equivalence

Key comparison BIPM.RI(II)-K1.Sn-113

MEASURAND: Equivalent activity of ^{113}Sn

Key comparison reference value: the SIR reference value x_{R} for this radionuclide is 58 840 kBq , with a standard uncertainty, u_{R} equal to 310 kBq (see Section 4.1 of the Final Report). The value x_i is taken as the equivalent activity for a laboratory i.

The degree of equivalence of each laboratory with respect to the reference value is given by a pair of terms: $D_i = (x_i - x_{\mathrm{R}})$ and U_i, its expanded uncertainty ($k = 2$), both expressed in kBq, and $U_i = 2((1 - 2w_i)u_i^2 + u_{\mathrm{R}}^2)^{1/2}$, where w_i is the weight of laboratory i contributing to the calculation of x_{R}.

Appendix B. Table of degrees of equivalence for BIPM.RI(II)-K1.Sn-113

Table B1: The table of degrees of equivalence for BIPM.RI(II)-K1.Sn-113

NMI i	D_i /**kBq**	U_i /**kBq**
PTB	300	1300
CIEMAT	-410	870
LNE-LNHB	-100	1200

Appendix C. Graph of degrees of equivalence with the KCRV for ^{113}Sn (as it appears in Appendix B of the MRA)

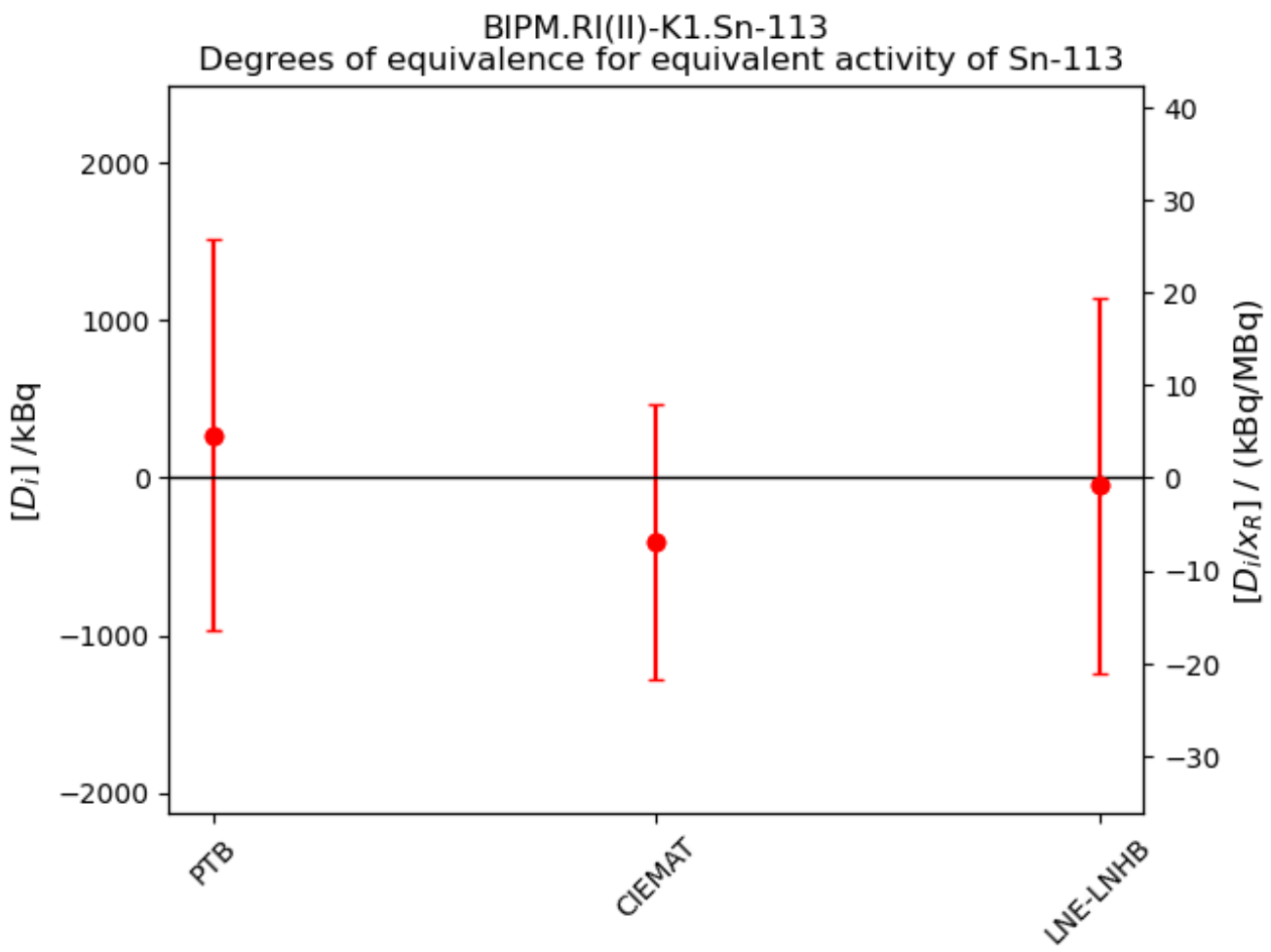

Figure C1. Degrees of equivalence for equivalent activity of ^{113}Sn.

Appendix D. Uncertainty budgets for the activity of ^{113}Sn submitted to the SIR

Detailed Uncertainty Budget

Laboratory: **LNE-LNHB**; Radionuclide: **^{113}Sn** ; Ampoule number: **3**

*Uncertainty components**, in % of the activity concentration, due to

Components	u / %	Comments	type	sensitivity factor
Counting statistics	**0.31**		**A**	1
Weighing	**0.016**		**B**	1
Source reproducibility	**0.5**		**A**	1
Dead time	**0**	Negligible		
Detection efficiency	**0.5**		**B**	1
Peak fitting model	**0.2**		**B**	1
Pile-up	**0.1**		**B**	1
Counting time	**0**	Negligible		
Adsorption	**0**	Not applicable		
Impurities	**0**	Not applicable		
Input parameters and statistical model	**0.26**	Photon emission intensity 391 keV	**B**	1
Decay correction	**0.01**		**B**	1
Combined uncertainty (as quadratic sum of all uncertainty component)	**0.85**			

* The uncertainty components are to be considered as approximations of the corresponding standard deviations (see also *Metrologia*, 1981, **17**, 73 and *Guide to expression of uncertainty in measurement*, ISO, corrected and reprinted 1995).

Appendix E. Acronyms used to identify different measurement methods

Each acronym has six components, geometry-detector (1)-radiation (1)-detector (2)-radiation (2)-mode. When a component is unknown, ?? is used and when it is not applicable 00 is used.

Geometry	**acronym**	**Detector**	**acronym**
4 π	4P	proportional counter	PC
defined solid angle	SA	press. Prop. Counter	PP
2 π	2P	liquid scintillation counting	LS
undefined solid angle	UA	NaI(Tl)	NA
		Ge(HP)	GH
		Ge(Li)	GL
		Si(Li)	SL
		CsI(Tl)	CS
		ionization chamber	IC
		grid ionization chamber	GC
		Cerenkov detector	CD
		calorimeter	CA
		solid plastic scintillator	SP
		PIPS detector	PS
		CeBr3	CB

Radiation	**acronym**	**Mode**	**acronym**
positron	PO	efficiency tracing	ET
beta particle	BP	internal gas counting	IG
Auger electron	AE	CIEMAT/NIST	CN
conversion electron	CE	sum counting	SC
mixed electrons	ME	coincidence	CO
bremsstrahlung	BS	anti-coincidence	AC
gamma rays	GR	coincidence counting with efficiency tracing	CT
x-rays	XR	anti-coincidence counting with efficiency tracing	AT
photons ($x + \gamma$)	PH	triple-to-double coincidence ratio counting	TD
photons + electrons	PE	selective sampling	SS
alpha particle	AP	high efficiency	HE
mixture of various radiation	MX	digital coincidence counting	DC

Examples of methods	**acronym**
4π(PC)β-γ coincidence counting	4P-PC-BP-NA-GR-CO
4π(PPC)β-γ coincidence counting eff. trac	4P-PP-MX-NA-GR-CT
defined solid angle α-particle counting with a PIPS detector	SA-PS-AP-00-00-00
4π(PPC)AX-γ(GeHP)-anticoincidence counting	4P-PP-MX-GH-GR-AC
4πCsI-β,AX,γ counting	4P-CS-MX-00-00-HE
calibrated IC	4P-IC-GR-00-00-00
internal gas counting	4P-PC-BP-00-00-IG