Update of the BIPM comparison BIPM.RI(II)-K1.Na-22 of activity measurements of the radionuclide ²²Na to include the 2014 result of the LNE-LNHB (France)

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Abstract Since 1976, 13 laboratories have submitted 29 samples of ²²Na 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.Na-22. Recently, the LNE-LNHB (France) participated in the comparison and the key comparison reference value (KCRV) has been updated. 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 $A_{\rm e}$, are all given in [1].

From its inception until 31 December 2019, the SIR has been used to measure 1016 ampoules to give 771 independent results for 72 different radionuclides. The SIR makes

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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 ongoing 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.Na-22 key comparison. The results of earlier participations in this key comparison were published previously [3–5].

2. Participants

Laboratory details are given in Table 1, with the earlier submissions being taken from [3–5]. 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.Na-22.

NMI or	Previous	Full name	Country	RMO	Date of
labora-	acronyms				SIR mea- surement
tory					yyyy-mm-dd
ANSTO	AAEC	Australian Nuclear Science	Australia	APMP	1981-09-10
		and Technology Organisa-			
		tion			
					1993-12-16
ASMW	-	Amt für Standardisierung,	former East	-	1977-11-24
		Meßwesen und Warenprü-	Germany		
		fung			
BARC	-	Bhabha Atomic Research	India	APMP	2004-07-13
		Centre			
BEV	IRK	Bundesamt fur Eich- und	Austria	EURAMET	2001-10-01
		Vermessungswesen			
BKFH	OMH,	Government Office of the	Hungary	EURAMET	1977-05-25
	MKEH	Capital City Budapest			
					1985-07-10
CIEMAT	-	Centro de Investigaciones	Spain	EURAMET	2006-05-12
		Energéticas, Medioambi-			
		entales y Tecnologicas			
CMI-IIR	UVVVR	Czech Metrological Insti-	Czech Re-	EURAMET	1977-05-13
		tute - Inspectorate for Ion-	public		
		izing Radiation			
			_		1979-05-15
LNE-	LMRI,	Laboratoire National de	France	EURAMET	1980-02-06
LNHB	LPRI	métrologie et d´Essais			
		-Laboratoire National			
		Henri Becquerel			1004.06.00
					1994-06-09
					2014-07-03

NMI or labora- tory	Previous acronyms	Full name	Country	RMO	Date of SIR measurement
NIST	NBS	National Institute of Stan-	United	SIM	1983-07-19
		dards and Technology	States		
NMIJ	ETL	National Metrology Insti- tute of Japan	Japan	APMP	1976-11-23
					1993-11-25
NMISA	NAC,	National Metrology Insti-	South Africa	AFRIMETS	1980-04-10
	CSIR-	tute of South Africa			
	NML ^a				
					1984-11-05
					2007-02-09
NPL	-	National Physical Labora-	United King-	EURAMET	1977-01-04
		tory	dom		
PTB	-	Physikalisch-Technische	Germany	EURAMET	1980-07-21
		Bundesanstalt			
					2006-05-22

... Continuation of Table 1.

3. NMI standardization methods

Each NMI that submits amoules 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 report [3–5]. The list of acronyms used to summarize the methods is given in Appendix E.

Since 2004, the half-life used by the BIPM is 950.8(9) days as published in IAEA TECDOC-619 [6]. The half-life of 950.4(15) days [7] was used for the earlier results.

^a NAC is another institute of the country.

Table 2: Standardization methods of the participants for $^{22}\mathrm{Na}.$

NMI or labora- tory	labora- acronym		$egin{array}{c} { m Relative} \ { m standard} \ { m uncertainty} \ /{ m 10^{-2}} \end{array}$		Reference date	Half-life /d
			A	В	yyyy-mm- dd	
ANSTO	$4\pi\beta$ - γ coincidence (4P-PC-PO-NA-GR-CO)	404.8	0.1	0.36	1981-08-15 00:00 UT	-
	$4\pi\beta$ - γ coincidence (4P-PC-PO-NA-GR-CO)	592.8	0.22	0.06	1991-03-17 23:00 UT	
ASMW	$4\pi\beta$ - γ coincidence (4P-PC-PO-NA-GR-CO)	2575.4 ^d	0.15	0.15	1977-10-21 12:00 UT	
		2602.4	0.15	0.15		
BARC	Ionization chamber calibrated traceable to NIST (4P-IC-GR-00-00)	1351	0.1	1.1	2004-02-16 06:30 UT	950.3602
	$4\pi\beta$ - γ coincidence (4P-PC-BP-NA-GR-CO)	1364	0.30	0.22		
BEV	Ionization chamber calibrated traceable to NPL (4P-IC-GR-00-00)	691.8	0.08	0.4	2001-09-01 00:00 UT	950.5
BKFH	$4\pi\beta$ - γ coincidence (4P-PC-PO-NA-GR-CO)	1384.3 ^d	0.2	0.6	1977-04-30 12:00 UT	950.4(15) [7]
		1384.3	0.2	0.6		
	$4\pi\beta$ - γ coincidence (4P-PC-	3626	0.04	0.36	1985-07-01	950.4(4)
CIEMAD	PO-NA-GR-CO)	0.40.4) 07	12:00 UT	[10]
CIEMAT	$4\pi\beta$ - γ coincidence and $4\pi\gamma$ NaI(Tl) well counter (4P-PP-PO-NA-GR-CO & 4P-NA-GR-00-00)	342.4	().37	2006-01-01 00:00 UT	950.6(4)
CMI-IIR	$4\pi\beta$ - γ coincidence (4P-PC-PO-NA-GR-CO)	3993	0.06	0.4	1977-04-14 13:00 UT	-
	$4\pi\beta$ - γ coincidence (4P-PC-PO-NA-GR-CO)	4198	0.2	0.4	1979-03-19 11:00 UT	949.7
LNE- LNHB	4π (NaI) γ counting (4P-NA-GR-00-00-00)	1765.8 ^d	0.01	0.2	1979-11-27 12:00 UT	950.4(4) [10]
		1747.5	0.01	0.2		
	$4\pi\beta$ - γ coincidence (4P-PC-PO-GH-GR-CO)	653.7 ^d	0.14	0.19	1994-05-18 12:00 UT	950.6(3)
	ĺ	652.3	0.14	0.19		
	Anti-coincidence counting (4P-PC-MX-NA-GR-AC)	1684.01	0.22	0.05	2014-06-16 12:00 UT	950.7(3)
	TDCR (4P-LS-MX-00-00- TD)	1684.01	0.13	0.27		
	$4\pi\gamma$ counting (4P-NA-GR-00-00-00)	1681.37	0.10	0.27		

... Continuation of Table 2.

NMI or	Method used and the	Activity	Relati	ve	Reference	Half-life
labora-	acronym	A_i/\mathbf{kBq}	standa		date	$/\mathbf{d}$
tory			uncert	ainty		
			$egin{array}{ c c c c c c c c c c c c c c c c c c c$	В		
			A		yyyy-mm- dd	
NIST	Ionization chamber cal-	3206	0.04	0.53	1983-06-06	-
	ibrated in 1969 using				14:00 UT	
	$4\pi\beta$ - γ coincidence and					
	annihilation- γ coincidence					
	(4P-PP-MX-NA-GR-CO &					
NINGER	4P-NA-GR-NA-GR-CO) $4\pi\beta$ -γ coincidence (4P-PC-	1897.4 ^d	0.02	0.28	1976-11-01	-
NMIJ	$4\pi\beta$ - γ coincidence (4P-PC-PO-NA-GR-CO)	1897.4	0.02	0.28	1976-11-01 12:00 UT	
	PO-NA-GR-CO)	1908.5	0.02	0.28	12:00 01	
	$4\pi\beta$ - γ coincidence (4P-PC-	1813	0.02	0.28	1993-11-01	
	PO-NA-GR-CO)	1010	0.4	0.5	00:00 UT	
NMISA	4π (liquid scintillation) β -	10 803 ^d	0.04	0.23	1980-03-18	950.4
111111111	γ coincidence (4P-LS-PO-	10000	0.01	0.20	10:00 UT	050.1
	NA-GR-CO) ^a					
	,	12 087	0.04	0.23		
	4π (liquid scintillation) β -	19 830	0.11	0.16	1984-10-24	
	γ coincidence (4P-LS-PO-				12:00 UT	
	NA-GR-CO) ^a					
	4π (liquid scintillation) β -	604.9	0.05	0.16	2006-08-16	950.6(4)
	γ coincidence (4P-LS-PO-				06:00 UT	[11]
	NA-GR-CO) ^b					
NPL	Ionization chamber cali-	591.6 ^d	0.15	1.98	1976-12-20	-
	brated using NPL primary				00:00 UT	
	standard (4P-IC-GR-00-00-					
	00)	200 0		1.00		
DED	4.2	592.8	0.15	1.98	1000 05 01	
PTB	$4\pi\beta$ - γ coincidence (4P-PC-	3622.1	0.04	0.1	1980-05-01	
	PO-NA-GR-CO) $4\pi\beta-\gamma \text{coincidence}^{c} \text{and}$	6779 ^{c2}	0.08	0.18	00:00 UT 2006-01-01	050 5(4)
	$4\pi\beta$ - γ coincidence ^c and CIEMAT-NIST (4P-	0779	0.08	0.18	00:00 UT	950.5(4)
	PC-PO-NA-GR-CO &				00.00 01	
	4P-LS-MX-00-00-CN) ^e					
	41 LB-WIX-00-00-01()					

^a see details in [8]

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.

b relying on the $\beta+$ branching ratio of 89.892 %

^c with distance variation of the NaI detector

^{c2} weighted mean of the two measurements taking into account correlation due to weighing (same balance)

^d Several samples submitted

^e see details in [9]

NMI or	Chemical	Solvent conc.	Carrier	Density	Relative activity of
laboratory	composi-		conc.		any impurity ^a
	tion				
/ SIR year		/ (mol dm ⁻³)	$/(\mu g g^{-1})$	$/(g \text{cm}^{-3})$	
ANSTO 1981	NaCl in HCl	0.1	NaCl: 200	1	<0.05 %
1993	²² Na in HCl	0.1	-	-	-
ASMW 1977	NaCl in HCl	0.1	NaCl: 20	-	<0.1 %
BARC 2004	NaCl in HCl	0.1	NaCl: 25	1	-
BEV 2001	NaCl in HCl	0.1	NaCl: 50	1	-
BKFH 1977	Na in HCl	0.1	Na: 25	-	<0.03 %
1985	Na in HCl	0.1	Na: 25	-	-
CIEMAT	NaCl in HCl	0.1	NaCl: 50	1	-
2006					
CMI-IIR	NaCl in wa-	-	NaCl:50	1	<0.1 %
1977	ter				
1979	NaCl in HCl	0.08	NaCl:50	-	<0.1 %
LNE-LNHB	NaCl in HCl	0.1	NaCl: 10	0.999	<0.0001 %
1980					
1994	NaCl in HCl	1	NaCl: 100	1.016	-
2014	NaCl in HCl	0.1	NaCl:10	1.0001	None detected
NIST 1983	NaCl in HCl	1	NaCl: 81	1.014	-
NMIJ 1976	NaCl in HCl	0.1	NaCl:50	-	-
1993	NaCl in HCl	0.1	NaCl:50	1	-
NMISA 1980	NaCl in HCl	1	Na ⁺ : 395	1.036	-
1984	NaCl in HCl	1	Na ⁺ :224	1.0169	-
2007	NaCl in HCl	1	Na ⁺ : 200	1.018	-
NPL 1977	NaCl in HCl	0.1	NaCl:100	-	<0.03 %
PTB 1980	NaCl in HCl	0.1	NaCl: 50	1	-
2006	NaCl in HCl	0.1	NaCl: 50	1	-

Table 3: Details of each solution of ²²Na submitted.

4. Results

All the submissions to the SIR since its inception in 1976 are maintained in a database known as the "master-file". The latest submission has added 1 ampoule for the activity measurements for 22 Na giving rise to 29 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]. No recent submission has been identified as a pilot study so the most recent result of each NMI is normally eligible for Appendix B of the MRA [2]. Since 1980, the LNE-LNHB participated to the SIR three times using three different primary

^a the ratio of the activity of the impurity to the activity of ²²Na at the reference date

measurement methods. It is worth noting that the three SIR comparison results obtained show a relative standard deviation of 4×10^{-4} .

Table 4: Results of SIR measurement of ²²Na.

NMI or labo-	m_i	A_i	$^{226}{ m Ra}$	$A_{\mathbf{e}i}$	Relative	$u_{\mathbf{c}i}$	$A_{\mathbf{e}i}$ for
ratory			source		uncert.		KCRV
-					from		
					SIR		
/ SIR year	$/\mathbf{g}$	$/\mathbf{k}\mathbf{B}\mathbf{q}$		/kBq	$/10^{-4}$	$/\mathbf{kBq}$	$/\mathbf{kBq}$
ANSTO 1981	3.596	404.8	3	7535	7	28	7535(28)
1993	3.728 27	592.8	2	7671	14 ^a	21] -
ASMW 1977	3.502 01	2575.4	4	7503	4	16	7502(16) ^f
	3.538 71	2602.4	4	7501	4	16	-
BARC 2004	3.635 34	1351	3	7506 ^b	5	80	7579(29)g
		1364		7579		29	-
BEV 2001	3.597 3	691.8	3	7490	6	31	-
BKFH 1977	3.605 8	1384.3	3	7547	6	44	-
	3.605 9	1384.3	3	7542	6	44	-
1985	3.603 9	3626	5	7525	4	27	7525(27)
CIEMAT 2006	3.645 8	342.4	2	7523	9	29	7523(29)
CMI-IIR 1977	3.530 96	3993	4	7526	5	31	-
1979	3.662 81	4198	4	7537	5	34	7537(34)
LNE-LNHB	3.610 99	1765.8	4	7528	5	16	-
1980							
	3.573 69	1747.5	4	7530	5	16	-
1994	3.567 4	653.7	3	7534	6	18	1 -
	3.559 4	652.3	3	7536	6	18	-
2014	3.777 5	1684.01	4	7533°	5	17	7533(17)
		1684.01		7533		23	_
		1681.37		7521		22	-
NIST 1983	3.725 58	3206	4	7496	3	40	7496(40)
NMIJ 1976	3.605 81	1897.4	4	7518	4	22	-
	3.626 86	1908.5	4	7516	4	22	-
1993	3.608 55	1813	4	7543	5	38	7543(38)
NMISA 1980	$3.594^{ m d}$	10803	5	7477	4	18	-
	3.610	12 087	5	7475	4	18	_
1984	3.619 52 ^e	19830	5	7527	4	15] -
2007	3.648 58	604.9	3	7568	6	14	7568(14)
NPL 1977	3.549 0	591.6	3	7538	5	150	$7540(150)^{f}$
	3.555 9	592.8	3	7542	5	150	_
PTB 1980	3.673 2	3622.1	4	7528	4	9	-
2006	3.600 44(9)	6779	5	7523	4	15	7523(15)

^a large SIR uncertainty due to the delay between the SIR measurement and the reference date

^b for the KCDB, the weighted mean activity and standard uncertainty are used to calculate the final result A_e =7570(27) kBq.

c result 7533(17) kBq from anti-coincidence counting used in KCDB

^d masses before dilution were 94.17 mg and 105.36 mg, respectively.

^e mass before dilution was 198.954 mg.

f An average value and average uncertainty between all submitted samples is used for the KCDB [12].

g Primary result

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 [13] 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 chisquared value is one. In addition, it allows for a power α smaller than two in the weighting factor. As proposed in [13], 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 recent result produces an updated KCRV for 22 Na in 2020 of **7534.0(73)** kBq with the power $\alpha = 1.75$ that has been calculated using the previously published results for the ASMW (1977), NPL (1977), CMI-IIR (1979), ANSTO (1981), NIST (1983), BKFH (1985), NMIJ (1993), BARC (2004) (primary result only), CIEMAT (2006), PTB (2006), NMISA (2007), and the present LNE-LNHB (2014) result. This can be compared with the previous KCRV values of 7526(5) kBq published

in 2003 [3] and 7534(7) kBq published in 2010 [5].

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). Normally, the most recent result is the one included. 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

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

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

$$U_i = 2u(D_i) (2)$$

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

$$u^{2}(D_{i}) = (1 - 2w_{i})u_{i}^{2} + u^{2}(KCRV)$$
(3)

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

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

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

$$u^{2}(D_{ij}) = u_{i}^{2} + u_{j}^{2} - 2u(A_{ei}, A_{ej})$$
(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 [14] 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 ongoing key comparison for ²²Na, BIPM.RI(II)-K1.Na-22 currently comprises 6 results. The KCRV has been recalculated including the result from LNE-LNHB (France) that superseeds their earlier participation in the comparison. The results have been analyzed with respect to the updated KCRV, providing degrees of equivalence for 6 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 ²²Na activity measurements to this comparison or take part in other linked comparisons.

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Appendix A. Introductory text for $^{22}\mathrm{Na}$ degrees of equivalence

Key comparison BIPM.RI(II)-K1.Na-22

MEASURAND: Equivalent activity of ²²Na

Key comparison reference value: the SIR reference value $x_{\rm R}$ for this radionuclide is 7534.0 kBq, with a standard uncertainty, $u_{\rm R}$ equal to 7.3 kBq (see Section 4.1 of the Final Report). The value x_i is taken as the equivalent activity for a laboratory i.

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

Appendix B. Table of degrees of equivalence for BIPM.RI(II)-K1.Na-22 $\,$

Table B1: The table of degrees of equivalence for ${\rm BIPM.RI(II)\text{-}K1.Na\text{-}22}$

NMI i	D_i / \mathbf{kBq}	U_i / \mathbf{kBq}
BEV	-44	64
BARC	36	56
CIEMAT	-11	56
PTB	-11	29
NMISA	34	27
LNE-LNHB	-1	32

Appendix C. Graph of degrees of equivalence with the KCRV for ²²Na (as it appears in Appendix B of the MRA)

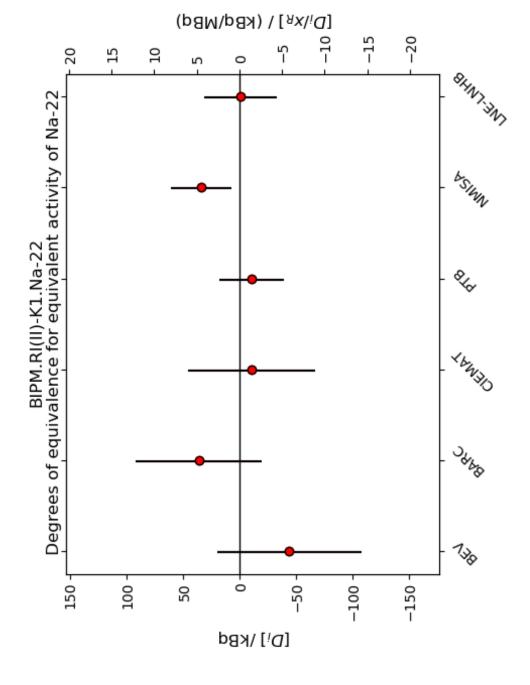


Figure C1. Degrees of equivalence for equivalent activity of $^{22}\mathrm{Na}$.

Appendix D. Uncertainty budgets for the activity of $^{22}\mathrm{Na}$ submitted to the SIR

Detailed Uncertainty Budget

Laboratory: LNE-LNHB;	Radionuclide: ²² N	a _ ; Ampoule number: 5	Method: anti	-coincidence
Uncertainty components*, in %	of the activity conc	entration, due to		
		Remarks	Evaluation	Relative
			type (A or B)	Sensitivity
				Factor
statistics	0,08	9 sources	A	
weighing	0,05	Pycnometer technique	В	
dead time	0,01	Live time technique	В	
background	0,1		A	
pile-up				
counting time				
adsorption				
impurities				
tracer input parameters				
and statistical model		/======================================		
quenching interpolation from	====			
calibration curve				
decay-scheme parameters half life ($T_{1/2}$: 2,6029 (8) a)	negligible		х В	
self absorption				
extrapolation of efficiency curve	0,18 I	Extrapolation technique	\mathbf{A}_{\perp}	
other effects (if relevant) (geometry factor, retrodiffusion)			x	
combined uncertainty (as quadratic sum of all uncertainty component	0,23			

^{*} 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).

Detailed Uncertainty Budget

Laboratory: LNE-LNHB; Radionuclide: ²²Na _ ; Ampoule number: 3 Method: TDCR

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

		Remarks	Evaluation	Relative
			type (A or B)	Sensitivity
				Factor
counting statistics	0,13	including sources variability	y A	1
weighing	0,10	pycnometer technique	В	1
dead time	0,10		В	1
background	0,01		Α	1
kB factor	0,02	uniform distribution	В	1
		[0,007-0,015] cm/MeV		
Compton spectrum	0,20	Monte Carlo simulation	В	1
decay scheme	0,10	P beta+	В	1
decay correction	0,001		В	1
combined uncertainty (as quadratic sum of all uncertainty components)	0,30	rounded		

^{*} 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).

Detailed Uncertainty Budget

Radionuclide: ²²Na; Ampoule number: 5 Method: 4π NaI well type counting Laboratory: LNE-LNHB; Uncertainty components*, in % of the activity concentration, due to **Evaluation** Relative Remarks type (A or B) Sensitivity Factor 0,1 Standard deviation of the statistics A mean 12 sources measured weighing 0,05 Pycnometer technique B dead time 0,01 Live-time technique В background 0,01 pile-up counting time adsorption impurities -----tracer ______ input parameters and statistical model quenching interpolation from calibration curve **Detection efficiency** 0,25 **Monte Carlo calculation** В decay-scheme parameters half life $(T_{1/2}: 2.6029 (8) a)$ negligible B

1981, 17, 73 and Guide to expression of uncertainty in measurement, ISO, corrected and reprinted 1995).

Conservative calculation

B

self absorption

zero-energy extrapolation

other effects (if relevant) (geometry factor, retrodiffusion)

combined uncertainty

(as quadratic sum of all

0,1

0,3

^{*} The uncertainty components are to be considered as approximations of the corresponding standard deviations (see also *Metrologia*,

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

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	СО
bremsstrahlung	BS	anti-coincidence	AC
gamma rays	GR	coincidence counting with	СТ
		efficiency tracing	
x-rays	XR	anti-coincidence counting	AT
		with efficiency tracing	
photons $(x + \gamma)$	PH	triple-to-double coincidence	TD
		ratio counting	
${ m photons} + { m electrons}$	PE	selective sampling	SS
alpha particle	AP	high efficiency	HE
mixture of various radi-	MX	digital coincidence counting	DC
ation			

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