



# Review Precise Half-Life Values for Two-Neutrino Double-β Decay: 2020 Review

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**Abstract:** All existing positive results on two-neutrino double beta decay and two-neutrino double electron capture in different nuclei have been analyzed. Weighted average and recommended half-life values for <sup>48</sup>Ca, <sup>76</sup>Ge, <sup>82</sup>Se, <sup>96</sup>Zr, <sup>100</sup>Mo, <sup>100</sup>Mo - <sup>100</sup>Ru (0<sub>1</sub><sup>+</sup>), <sup>116</sup>Cd, <sup>128</sup>Te, <sup>130</sup>Te, <sup>136</sup>Xe, <sup>150</sup>Nd, <sup>150</sup>Nd - <sup>150</sup>Sm (0<sub>1</sub><sup>+</sup>), <sup>238</sup>U, <sup>78</sup>Kr, <sup>124</sup>Xe and <sup>130</sup>Ba have been obtained. Given the measured half-life values, effective nuclear matrix elements for all these transitions were calculated.

**Keywords:** double beta decay; nuclear matrix elements; <sup>48</sup>Ca; <sup>76</sup>Ge; <sup>82</sup>Se; <sup>96</sup>Zr; <sup>100</sup>Mo; <sup>100</sup>Mo - <sup>100</sup>Ru (0<sub>1</sub><sup>+</sup>); <sup>116</sup>Cd; <sup>128</sup>Te; <sup>130</sup>Te; <sup>136</sup>Xe; <sup>150</sup>Nd; <sup>150</sup>Nd - <sup>150</sup>Sm (0<sub>1</sub><sup>+</sup>); <sup>238</sup>U; <sup>78</sup>Kr; <sup>124</sup>Xe; <sup>130</sup>Ba

## 1. Introduction

Two-neutrino double beta decay  $(2\nu\beta\beta)$  was first considered by Maria Goeppert-Mayer in 1935 [1]:

$$(A, Z) \to (A, Z+2) + 2e^- + 2\bar{\nu}$$
 (1)

This is a process in which a nucleus (A,Z) decays to a nucleus (A,Z + 2) by emitting two electrons and two electron-type antineutrinos. The  $2\nu\beta\beta$  decay is a second-order weak interaction process and does not violate any conservation laws. Nevertheless, the study of this process provides rich information that can be used both to clarify various aspects of neutrinoless double beta decay and to search for exotic processes (decays with Majoron emission [2,3], bosonic neutrinos [4], violation of Lorentz invariance [3,5,6], the presence of right-handed leptonic currents [7], neutrino self-interactions ( $\nu$ SI) [8], etc.). The  $2\nu\beta\beta$  decay was first discovered in a geochemical experiment with <sup>130</sup>Te in 1950 [9]. In a direct (counter) experiment, the decay was first recorded by M. Moe et al. in 1987 (TRC, <sup>82</sup>Se) [10]. To date,  $2\nu\beta\beta$  decay has already been studied quite well. This process has been registered for 11 nuclei. For some nuclei (<sup>100</sup>Mo, <sup>150</sup>Nd), a transition to the 0<sup>+</sup><sub>1</sub> excited state of the daughter nucleus was detected too. In addition, a two-neutrino double electron capture (ECEC(2 $\nu$ )) was detected in several nuclei (<sup>130</sup>Ba [11], <sup>124</sup>Xe [12], <sup>78</sup>Kr [13]). In this process, two orbital electrons are captured. In the final state, two neutrinos and two X-rays appear:

$$e^- + e^- + (A, Z) \to (A, Z - 2) + 2\nu + 2X.$$
 (2)

In the NEMO-3 experiment, all decay characteristics (total energy spectrum, single electron spectrum, angular distribution) for 7 isotopes (<sup>48</sup>Ca, <sup>82</sup>Se, <sup>96</sup>Zr, <sup>100</sup>Mo, <sup>116</sup>Cd, <sup>130</sup>Te and <sup>150</sup>Nd) were studied simultaneously. At present, the study of two-neutrino processes is moving into a new stage, precision study. The accuracy of determining the half-life values and other characteristics of this process is becoming increasingly important (see the discussion in [3,7,14–16]). The exact half-lives are important to know for the following reasons:

1. Nuclear spectroscopy. It has now been established that some isotopes that were previously considered stable are not, and decay of these isotopes is observed through the  $2\nu\beta\beta$  decay with a

half-life of  $\sim 10^{18} - 10^{24}$  yr. One just need to know the exact half-life values to include them in the isotope tables. Then, these values can be used for any purpose.

2. Nuclear matrix elements (NME). First of all, one can check the quality of NME calculations for  $2\nu\beta\beta$  decay, because it is possible to directly compare experimental and calculated values. Secondly, accurate knowledge of the NME( $2\nu$ ) also makes it possible to improve the quality of NME calculations for neutrinoless double beta decay ( $0\nu\beta\beta$ ). For example, the accurate half-life values for  $2\nu\beta\beta$  decay are used to determine the most important parameter of the quasiparticle random-phase approximation model (QRPA), the strength of the particle–particle interaction  $g_{pp}$  [17–19].

3. To fix  $g_A$  (axial-vector coupling constant). There are indications that, in nuclear medium, the matrix elements of the axial-vector operator are reduced in comparison with their free nucleon values. This quenching is described as a reduction of the coupling constant  $g_A$  from its free nucleon value of  $g_A = 1.2701$  [20] to the value of  $g_A \sim 0.35$ -1.0 (see [21–23]). In principle,  $g_A$  value could be established by comparison of exact experimental values and results of theoretical calculations of NMEs. Finally, it can help in understanding the  $g_A$  value in the case of  $0\nu\beta\beta$  decay (see discussions in Refs. [21–23]).

It should be noted here that the phenomenological interpretation of the change in the value of  $g_A$  in nuclear matter is apparently connected with the imperfection of our description of the nuclear structure and the process of double beta decay itself. Therefore, when describing the process of  $2\nu\beta\beta$  decay, we are essentially adjusting the value of  $g_A$  in order to give a correct description of the process. In this sense, this is about the same as the situation with  $g_{pp}$  in the previous paragraph.

4. A check of the "bosonic" neutrino hypothesis [4] and  $\nu$ SI [8].

At the same time, it is quite difficult to choose the "best" result from the available data. For some isotopes, up to 7–10 different measurements exist. The quality of these results is not always obvious. Therefore, it is difficult to choose the best ("correct") value for the half-life.

In the present paper, a critical analysis of all available results on two-neutrino decay has been performed and average or/and recommended half-life values for all isotopes are presented. Using these values and the values of the phase space factors from [24,25], the "effective" NMEs were calculated.

The first time that such type of work was done was in 2001, and the results were presented at the International Workshop on the calculation of double beta decay nuclear matrix elements, MEDEX'01 [26]. Then, updated half-life values were presented at MEDEX'05, MEDEX'09 and MEDEX'13 and published in Refs. [14,15,27], respectively. In this article, new positive results obtained since the beginning of 2015 and to the middle of 2020 have been added and analyzed. Preliminary results of this analysis have been presented at MEDEX'19 [28].

The main differences from the previous analysis [15] are the following: (1) The new experimental results are included in the analysis:  ${}^{48}$ Ca [29],  ${}^{76}$ Ge [30],  ${}^{82}$ Se [31,32],  ${}^{100}$ Mo [3,33],  ${}^{116}$ Cd [34,35],  ${}^{130}$ Te [36,37],  ${}^{136}$ Xe [38],  ${}^{150}$ Nd [39],  ${}^{150}$ Nd -  ${}^{150}$ Sm (0<sup>+</sup><sub>1</sub>) [40] and  ${}^{130}$ Ba [41]; (2) the positive results obtained for  ${}^{78}$ Kr [13] and  ${}^{124}$ Xe [12] are added (these decays have been detected for the first time). I would like to stress that most of the above-mentioned new results are very precise. The accuracy of some of the obtained half-life values is ~2–3%. This is a result of (mainly) experiments with low-temperature bolometers (including scintillating bolometers).

#### 2. Experimental Data

Tables 1 and 2 show the experimental results on  $2\nu\beta\beta$  decay and on ECEC( $2\nu$ ) capture in different nuclei. For direct experiments, the number of detected (useful) events and the signal-to-background (S/B) ratio are presented.

Nucleus	Ν	<i>T</i> <sub>1/2</sub> , yr	S/B	Ref., Year
<sup>48</sup> Ca	$\sim 100$	$[4.3^{+2.4}_{-1.1}(stat) \pm 1.4(syst)] \cdot 10^{19}$	1/5	[42], 1996
	5	$4.2^{+3.3}_{-1.3} \cdot 10^{19}$	5/0	[43], 2000
	116	$\frac{4.2^{+3.3}_{-1.3} \cdot 10^{19}}{[6.4^{+0.7}_{-0.6}(stat)^{+1.2}_{-0.9}(syst)] \cdot 10^{19}}$	3.9	[ <mark>29</mark> ], 2016
		Average value: $5.3^{+1.2}_{-0.8} \cdot 10^{19}$		
<sup>76</sup> Ge	$\sim 4000$	$(0.9\pm 0.1)\cdot 10^{21}$	$\sim 1/8$	[44], 1990
	758	$1.1^{+0.6}_{-0.3} \cdot 10^{21}$	$\sim 1/6$	[ <b>45</b> ], 1993
	$\sim 330$	$\begin{array}{c} 1.1^{+0.6}_{-0.3} \cdot 10^{21} \\ 0.92^{+0.07}_{-0.04} \cdot 10^{21} \\ \end{array}$	$\sim 1.2$	[ <b>46</b> ], 1993
	132	$1.27_{-0.16}^{+0.21} \cdot 10^{21}$	$\sim 1.4$	[47], 1994
	$\sim 3000$	$(1.45 \pm 0.15) \cdot 10^{21}$	$\sim 1.5$	[ <b>48</b> ], 1999
	$\sim \! 80,\! 000$	$[1.74 \pm 0.01(stat)^{+0.18}_{-0.16}(syst)] \cdot 10^{21}$	$\sim 1.5$	[49], 2003
	25,690	$(1.925 \pm 0.094) \cdot 10^{21}$	$\sim 3$	[ <mark>30</mark> ], 2015
		Average value: $(1.88\pm0.08)\cdot10^{21}$		
<sup>82</sup> Se	89.6	$1.08^{+0.26}_{-0.06} \cdot 10^{20}$	$\sim 8$	[ <mark>50</mark> ], 1992
	149.1	$[0.83 \pm 0.10(stat) \pm 0.07(syst)] \cdot 10^{20}$	2.3	[ <mark>51</mark> ], 1998
	2750	$[0.939 \pm 0.017(stat) \pm 0.058(syst)] \cdot 10^{20}  ^{(a)}$	4	[31], 2018
	$\sim 200,000$	$[0.860 \pm 0.003 (stat) ^{+0.019}_{-0.013} (syst)] \cdot 10^{20} \ ^{(a)}$	$\sim 10$	[ <mark>32</mark> ], 2019
		$(1.3 \pm 0.05) \cdot 10^{20}$ (geochem.)		[52], 198
		Average value: $0.87^{+0.02}_{-0.01} \cdot 10^{20}$		
<sup>96</sup> Zr	26.7	$[2.1^{+0.8}_{-0.4}(stat) \pm 0.2(syst)] \cdot 10^{19}$	$1.9^{(b)}$	[53], 1999
	453	$[2.35 \pm 0.14(stat) \pm 0.16(syst)] \cdot 10^{19}$	1	[54], 2010
		$(3.9 \pm 0.9) \cdot 10^{19}$ (geochem.)		[55], 1993
		$(0.94 \pm 0.32) \cdot 10^{19}$ (geochem.)		[56], 2003
		Average value: $(2.3\pm0.2)\cdot10^{19}$		
<sup>100</sup> Mo	$\sim 500$	$11.5^{+3.0}_{-2.0}\cdot 10^{18}$	1/7	[57], 1991
	67	$11.6^{+\overline{3.4}}_{-0.8}\cdot 10^{18}$	7	[ <mark>58</mark> ], 1993
	1433	$[7.3 \pm 0.35(stat) \pm 0.8(syst)] \cdot 10^{18} \ ^{(a)(c)}$	3	[ <mark>59</mark> ], 1995
	175	$7.6^{+2.2}_{-1.4} \cdot 10^{18}$	1/2	[ <mark>60</mark> ], 1992
	377	$[6.82^{+0.38}_{-0.53}(stat) \pm 0.68(syst)] \cdot 10^{18}$	10	[ <mark>61</mark> ], 1997
	800	$[7.2 \pm 1.1(stat) \pm 1.8(syst)] \cdot 10^{18}$	1/9	[ <mark>62</mark> ], 2001
	$\sim 350$	$[7.15 \pm 0.37(stat) \pm 0.66(syst)] \cdot 10^{18}$	$\sim 5^{~(d)}$	[ <mark>63</mark> ], 2014
	500,000	$[6.81 \pm 0.01 (stat) ^{+0.38}_{-0.40} (syst)] \cdot 10^{18} \ ^{(a)}$	80	[3], 2019
	35,638	$[7.12^{+0.18}_{-0.14}(stat) \pm 0.10(syst)] \cdot 10^{18}$ <sup>(a)</sup>	10	[ <mark>33</mark> ], 2020
		$(2.1 \pm 0.3) \cdot 10^{18}$ (geochem.)		[ <mark>64</mark> ], 2004
		Average value: $7.06^{+0.15}_{-0.13} \cdot 10^{18}$		
<sup>100</sup> Mo -	133 <sup>(e)</sup>	$6.1^{+1.8}_{-1.1} \cdot 10^{20}$	1/7	[ <b>65</b> ], 1995
$^{100}$ Ru (0 <sup>+</sup> <sub>1</sub> )	153 <sup>(e)</sup>	$[9.3^{+2.8}_{-1.7}(stat) \pm 1.4(syst)] \cdot 10^{20}$	1/4	[66], 1999
	19.5	$[5.9^{+1.7}_{-1.1}(stat) \pm 0.6(syst)] \cdot 10^{20}$ $[5.9^{+1.7}_{-1.1}(stat) \pm 0.6(syst)] \cdot 10^{20}$	${\sim}8$	[ <mark>67</mark> ], 2001
	35.5	$\begin{array}{l} [5.5\substack{+1.2\\-0.8}(stat)\pm0.3(syst)]\cdot10^{20}\\ [5.7\substack{+1.3\\-0.9}(stat)\pm0.8(syst)]\cdot10^{20}\end{array}$	${\sim}8$	[68], 2009
	37.5	$[5.7^{+1.3}_{-0.9}(stat) \pm 0.8(syst)] \cdot 10^{20}$	$\sim 3$	[ <mark>69</mark> ], 2007
	597 <sup>(e)</sup>	$[6.9^{+1.0}_{-0.8}(stat) \pm 0.7(syst)] \cdot 10^{20}$	$\sim 1/10$	[70], 2010
	239 <sup>(e)</sup>	$[7.5 \pm 0.6(stat) \pm 0.6(syst)] \cdot 10^{20}$	2	<b>[71], 2</b> 014
		Average value: $6.7^{+0.5}_{-0.4} \cdot 10^{20}$		
<sup>116</sup> Cd	$\sim 180$	$2.6^{+0.9}_{-0.5} \cdot 10^{19}$	$\sim 1/4$	[ <b>72</b> ], 1995
	174.6	$[2.9 \pm 0.3(stat) \pm 0.2(syst)] \cdot 10^{19} \ ^{(a)(c)}$	3	[ <b>73</b> ], 1996
	9850	$[2.9 \pm 0.06(stat)^{+0.4}_{-0.3}(syst)] \cdot 10^{19}$	$\sim 3$	[74], 2003
	4968	$[2.74 \pm 0.04(stat) \pm 0.18(syst)] \cdot 10^{19}$ (a)	12	[ <mark>34</mark> ], 2012
	93,000	$2.63^{+0.11}_{-0.12} \cdot 10^{19}$	1.5	[35], 2018
		Average value: $(2.69\pm0.09)\cdot10^{19}$		
<sup>128</sup> Te		$\sim 2.2 \cdot 10^{24}$ (geochem.)		[75], 1993
		$(7.7 \pm 0.4) \cdot 10^{24}$ (geochem.)		[76], 1993
		$(2.41 \pm 0.39) \cdot 10^{24}$ (geochem.)		[77], 2008
		$(2.3 \pm 0.3) \cdot 10^{24}$ (geochem.)		

**Table 1.** Present, positive  $2\nu\beta\beta$  decay results. N is the number of useful events, S/B is the signal-to-background ratio.

Nucleus	Ν	<i>T</i> <sub>1/2</sub> , yr	S/B	Ref., Year
		Recommended value: $(2.25\pm0.09)\cdot10^{24}\text{(f)}$		
<sup>130</sup> Te	260	$[6.1 \pm 1.4(stat)^{+2.9}_{-3.5}(syst)] \cdot 10^{20}$	1/8	[79], 2003
	236	$[7.0 \pm 0.9(stat) \pm 1.1(syst)] \cdot 10^{20}$	1/3	[ <mark>80</mark> ], 2011
	~33,000	$[8.2 \pm 0.2(stat) \pm 0.6(syst)] \cdot 10^{20}$	0.1-0.5	[36], 2017
	$\sim 20,000$	$[7.9 \pm 0.1(stat) \pm 0.2(syst)] \cdot 10^{20}$	>1	[37], 2020
		$\sim 8\cdot 10^{20}$ (geochem.)		[75], 1991
		$(27 \pm 1) \cdot 10^{20}$ (geochem.)		[ <mark>76</mark> ], 1993
		$(9.0 \pm 1.4) \cdot 10^{20}$ (geochem.)		[77], 2008
		$(8.0 \pm 1.1) \cdot 10^{20}$ (geochem.)		[78], 2008
		Average value: $(7.91\pm0.21)\cdot10^{20}$		
<sup>136</sup> Xe	~19,000	$[2.165 \pm 0.016(stat) \pm 0.059(syst)] \cdot 10^{21}$	$\sim 10$	[81], 2014
	$\sim \! 100,\! 000$	$[2.21 \pm 0.02(stat) \pm 0.07(syst)] \cdot 10^{21}$	$\sim \! 10$	[38], 2016
		Average value: $(2.18\pm0.05)\cdot10^{21}$		
<sup>150</sup> Nd	23	$[18.8^{+6.9}_{-3.9}(stat) \pm 1.9(syst)] \cdot 10^{18}$	1.8	[ <mark>82</mark> ], 1995
	414	$[6.75^{+0.37}_{-0.42}(stat) \pm 0.68(syst)] \cdot 10^{18}$	6	[ <mark>61</mark> ], 1997
	2214	$[9.34 \pm 0.22(stat)^{+0.62}_{-0.60}(syst)] \cdot 10^{18}$	4	[39], 2016
		Average value: $(8.4\pm1.1)\cdot10^{18}$		
		Recommended value: $(9.34\pm0.65)\cdot10^{18}$		
<sup>150</sup> Nd -	177.5 <sup>(e)</sup>	$[1.33^{+0.36}_{-0.23}(stat)^{+0.27}_{-0.13}(syst)] \cdot 10^{20}$	1/5	[83], 2009
$^{150}$ Sm (0 <sup>+</sup> <sub>1</sub> )	21.6	$[1.07^{+0.45}_{-0.25}(stat) \pm +0.07(syst)] \cdot 10^{20}$	$\sim 1.2$	[84], 2014
、 1 <i>'</i>	$\sim 6$	$[0.69^{+0.40}_{-0.19}(stat) \pm +0.11(syst)] \cdot 10^{20}$	$\sim 2$	[40], 2019
		Average value: $1.2^{+0.3}_{-0.2} \cdot 10^{20}$		
<sup>238</sup> U		$(2.0 \pm 0.6) \cdot 10^{21}$ (radiochem.)		[85], 1993

Table 1. Cont.

<sup>(a)</sup> For SSD mechanism. <sup>(b)</sup> For  $E_{2e} > 1.2$  MeV. <sup>(c)</sup> After correction (see [14]). <sup>(d)</sup> For  $E_{2e} > 1.5$  MeV. <sup>(e)</sup> In both peaks. <sup>(f)</sup> This value was obtained using average  $T_{1/2}$  for <sup>130</sup>Te and well-known ratio  $T_{1/2}(^{130}\text{Te})/T_{1/2}(^{128}\text{Te}) = (3.52 \pm 0.11) \cdot 10^{-4}$  [76].

**Table 2.** Present, positive two-neutrino double electron capture results. N is the number of useful events, S/B is the signal-to-background ratio. In the case of <sup>78</sup>Kr and <sup>124</sup>Xe  $T_{1/2}$  for  $2K(2\nu)$ , capture is presented (this is ~75–80% of  $ECEC(2\nu)$ ).

Nucleus	Ν	$T_{1/2}(2\nu)$ , yr		Ref., Year	
<sup>130</sup> Ba		$2.1^{+3.0}_{-0.8} \cdot 10^{21}$ (geochem.)		[ <mark>86</mark> ], 1996	
$ECEC(2\nu)$		$(2.2 \pm 0.5) \cdot 10^{21}$ (geochem.)		[ <b>11</b> ], 2001	
		$(0.60 \pm 0.11) \cdot 10^{21}$ (geochem.)		[87], 2009	
Recommended value: $(2.2 \pm 0.5) \cdot 10^{21}$					
<sup>78</sup> Kr	15	$[1.9^{+1.3}_{-0.7}(stat) \pm 0.3(syst)] \cdot 10^{22}$	15	[13], 2017	
$2K(2\nu)$					
		Recommended value: $(1.9^{+1.3}_{-0.8}) \cdot 10^{22}$ (?) $^{(a)}$			
<sup>124</sup> Xe	126	$[1.8 \pm 0.5(stat) \pm 0.1(syst)] \cdot 10^{22}$	0.2	[12], 2019	
$2K(2\nu)$					
Recommended value: $(1.8 \pm 0.5) \cdot 10^{22}$					
		(a) Soo toxt			

<sup>(a)</sup> See text.

### 3. Data Analysis

To calculate an average of the ensemble of available data, a standard procedure, as recommended by the Particle Data Group [20], was used. The weighted average and the corresponding error were calculated, as follows:

$$\bar{x} \pm \delta \bar{x} = \sum w_i x_i / \sum w_i \pm (\sum w_i)^{-1/2},$$
 (3)

where  $w_i = 1/(\delta x_i)^2$ . Here,  $x_i$  and  $\delta x_i$  are the value and error reported by the *i*-th experiment, and the summations run over the N experiments.

Then, it is necessary to calculate  $\chi^2 = \sum w_i(\bar{x} - x_i)^2$  and compare it with N - 1, which is the expectation value of  $\chi^2$  if the measurements are from a Gaussian distribution. In the case when  $\chi^2/(N-1)$  is less than or equal to 1 and there are no known problems with the data, then one accepts the results. In the case when  $\chi^2/(N-1) >> 1$ , one chooses not to use the average procedure at all. Finally, if  $\chi^2/(N-1)$  is larger than 1, but not greatly so, it is still best to use the average data, but to increase the quoted error,  $\delta \bar{x}$  in Equation (1), by a factor of S defined by

$$S = [\chi^2 / (N-1)]^{1/2}.$$
 (4)

For averages, the statistical and systematic errors are treated in quadrature and used as a combined error  $\delta x_i$ . In some cases, only the results obtained with a high enough S/B ratio were used.

## 3.1. <sup>48</sup>Ca

The  $2\nu\beta\beta$  decay of <sup>48</sup>Ca was observed in three independent experiments [29,42,43]. The obtained results are in good agreement. The weighted average value is:

$$T_{1/2} = 5.3^{+1.2}_{-0.8} \cdot 10^{19} \text{ yr}.$$

This value is slightly higher than the average value obtained in previous analysis  $(T_{1/2} = 4.4^{+0.6}_{-0.5} \cdot 10^{19} \text{ yr [15]})$ . This is due to the fact that the final result of the NEMO-3 experiment [29] was used in present analysis (the intermediate result of the NEMO-3 experiment [88] was used in [15]). The change in the final result in the NEMO-3 experiment was mainly due to the fact that after disassembling the detector, the parameters of sources containing <sup>48</sup>Ca were refined. It was found that, in reality, the diameter of the sources turned out to be slightly larger (and the thickness, respectively, less) than previously assumed. Taking this circumstance into account led to an increase in the calculated efficiency of recording useful events and, ultimately, to an increase in the  $T_{1/2}$  value for <sup>48</sup>Ca. In addition, systematic error in [29] is higher then in [88].

## 3.2. <sup>76</sup>Ge

For <sup>76</sup>Ge, a lot of positive results were obtained, but the scatter of the obtained values is rather large. Half-life values gradually increased over time during the 90-th. It was decided not to use the results of the early works (1990s), as a recent historical review [89] emphasized that the contribution of background processes was underestimated in these works. Therefore, to determine the average value, the results published after 2000 have been used, with large statistics and a high S/B ratio [30,49]. Note that the final result of the Heidelberg–Moscow collaboration was used [49]. As a result, we get:

$$T_{1/2} = (1.88 \pm 0.08) \cdot 10^{21} \text{ yr.}$$

#### 3.3. <sup>82</sup>Se

There are many geochemical measurements (~20) and only four independent counting experiments for <sup>82</sup>Se. However, the geochemical results are in poor agreement with each other and with the results of direct experiments. It is known that the possibility of existing large systematic errors in geochemical measurements cannot be excluded (see discussion in Ref. [90]). Thus, only the results of the direct measurements [31,32,50,51] were used to obtain a present half-life value for <sup>82</sup>Se. Single State Dominance (SSD) mechanism (see explanation in [91]) was established for  $2\nu\beta\beta$  transition in <sup>82</sup>Se [31,32] and half-life values in this papers were obtained under the assumption of

the SSD mechanism <sup>1</sup>. The result of Ref. [10] has not been used in the analysis because this is the preliminary result of [50]. The result of work [50] is presented with very asymmetrical errors. To be more conservative, the value for the lower error was taken to be the same as the upper one in our analysis. Finally, the weighted average value is:

$$T_{1/2} = 0.87^{+0.02}_{-0.01} \cdot 10^{20} \text{ yr}.$$

3.4. <sup>96</sup>Zr

There are two positive results from the direct experiments (NEMO-2 [53] and NEMO-3 [54]) and two geochemical results [55,56]. Taking into account the comment in Section 3.3, the values from direct experiments (Refs. [53,54]) were used to obtain a present weighted half-life value for <sup>96</sup>Zr:

$$T_{1/2} = (2.3 \pm 0.2) \cdot 10^{19} \,\mathrm{yr}$$

3.5. <sup>100</sup>Mo

By the present nine positive results from direct experiments<sup>2</sup> and one result from a geochemical experiment have been obtained. I do not use the geochemical result here (see comment in Section 3.3). Finally, in calculating the average, only the results of experiments with S/B ratios greater than 1 were used (i.e., the results of Refs. [3,33,59,61,63]). I use only final result of Elliott et al. [61] and do not consider their preliminary result from [58]. For <sup>100</sup>Mo SSD mechanism was installed and in Ref. [3,33,59] the half-lives were obtained taking this fact into account. In addition, the corrected half-life value from Ref. [59] has been used (see explanation in [14]). The following weighted average value for the half-life is obtained as:

$$T_{1/2} = 7.06^{+0.15}_{-0.13} \cdot 10^{18} \text{ yr}$$

3.6.  ${}^{100}Mo - {}^{100}Ru (0^+_1; 1130.32 \text{ Kev})$ 

The  $2\nu\beta\beta$  decay of <sup>100</sup>Mo to the 0<sup>+</sup><sub>1</sub> excited state of <sup>100</sup>Ru was detected in seven independent experiments. The results are in good agreement. The weighted average value for the half-life has been obtained using the results from [65,66,68–71]:

$$T_{1/2} = 6.7^{+0.5}_{-0.4} \cdot 10^{20} \text{ yr.}$$

The result from [68] was used as the final result of the TUNL-ITEP experiment (the result from [67] was not used here because I consider it as preliminary one).

3.7. <sup>116</sup>Cd

Five independent positive results were obtained [34,35,72–74]. The results are in good agreement with each other. The corrected result for the half-life value from Ref. [73] is used here. The original

<sup>&</sup>lt;sup>1</sup> It was experimentally demonstrated that in some nuclei ( $^{82}$ Se,  $^{100}$ Mo and  $^{116}$ Cd) the SSD mechanism is realized. In this case, the spectra (total energy, single electron energy and angular distribution) differ from the case of the High State Dominance (HSD) mechanism. In principle, this does not affect the half-life of the corresponding nuclei. In a real experiment, energy is recorded with a certain threshold, which can affect the efficiency of recording useful events. The neglect of this effect can lead to an error in the determination of  $T_{1/2}$  (up to ~ 10–15%). This is especially noticeable in experiments where the energy of an individual electron is recorded (for example, the NEMO-3 experiment).

<sup>&</sup>lt;sup>2</sup> I do not consider here the result of Ref. [92] because of a high background contribution that was not excluded in this experiment. As a result, the "positive" effect is mainly associated with the background. Calculations show that without the background contribution to the "positive" effect, the sensitivity of the experiment was simply not enough to detect <sup>100</sup>Mo decay.

half-life value was decreased by  $\sim 25\%$  (see explanation in [14]). In Refs. [34,73], half-life values were obtained with the assumption that the SSD mechanism was realized. The weighted average value is:

$$T_{1/2} = (2.69 \pm 0.09) \cdot 10^{19} \,\mathrm{yr}$$

### 3.8. <sup>128</sup>Te and <sup>130</sup>Te

There are a large number of geochemical results for these isotopes. Although the half-life ratio for these isotopes is well known (accuracy is ~3% [76]), the absolute  $T_{1/2}$  values for each isotope are different from one experiment to the next. One group of authors [75,93,94] gives  $T_{1/2} \approx 0.8 \cdot 10^{21}$  yr for <sup>130</sup>Te and  $T_{1/2} \approx 2 \cdot 10^{24}$  yr for <sup>128</sup>Te, while another group [52,76] claims  $T_{1/2} \approx (2.5 - 2.7) \cdot 10^{21}$  yr and  $T_{1/2} \approx 7.7 \cdot 10^{24}$  yr, respectively. In addition, as a rule, experiments with young samples (~100 million years) give results of the half-life value for <sup>130</sup>Te in the range of ~  $(0.7 - 0.9) \cdot 10^{21}$  yr, while experiments with old samples (> 1 billion years) give half-life values in the range of ~  $(2.5 - 2.7) \cdot 10^{21}$  yr. In 2008, it was demonstrated that short half-lives are more likely to be correct [77,78]. In a new experiment with young minerals, the half-life values were estimated at  $(9.0 \pm 1.4) \cdot 10^{20}$  yr [77] and  $(8.0 \pm 1.1) \cdot 10^{20}$  yr [78] for <sup>130</sup>Te and  $(2.41 \pm 0.39) \cdot 10^{24}$  yr [77] and  $(2.3 \pm 0.3) \cdot 10^{24}$  yr [78] for <sup>128</sup>Te. In fact, in both experiments, the half-life was measured only for <sup>130</sup>Te, and the value for <sup>128</sup>Te was determined using the previously measured  $T_{1/2}(^{130}\text{Te})/T_{1/2}(^{128}\text{Te})$  ratio [76]. If we average the values obtained in these two experiments, we get:  $T_{1/2} = (8.4 \pm 0.9) \cdot 10^{20}$  years for <sup>130</sup>Te and  $T_{1/2} = (2.3 \pm 0.3) \cdot 10^{24}$  yr eas for <sup>128</sup>Te, which is in good agreement with the results of direct (counter) experiments (see below).

The first indication of the observation of the  $2\nu\beta\beta$  decay for <sup>130</sup>Te in a direct experiment was obtained in [79]. More accurate and reliable values were obtained later in the NEMO-3 experiment [80]. Very precise results were obtained recently in CUORE-0 [36] and CUORE [37] experiments. The results are in good agreement, and the weighted average value is

$$T_{1/2} = (7.91 \pm 0.21) \cdot 10^{20} \text{ yr}$$

Now, using the very well-known ratio  $T_{1/2}(^{130}\text{Te})/T_{1/2}(^{128}\text{Te}) = (3.52 \pm 0.11) \cdot 10^{-4}$  [76], one can obtain the half-life value for  $^{128}\text{Te}$ ,

$$T_{1/2} = (2.25 \pm 0.09) \cdot 10^{24} \text{ yr}$$

I recommend using these two results as the most correct and reliable half-life values for <sup>130</sup>Te and <sup>128</sup>Te. As one can see now, results of direct and geochemical experiments are in good agreement.

3.9. <sup>136</sup>Xe

The half-life value for  $^{136}Xe$  was measured in two independent experiments, EXO [81,95,96] and Kamland-Zen [38,97,98]. To obtain the average value of the half-life, the most accurate results of these experiments obtained in [38,81] were used (see Table 1). The weighted average value is

$$T_{1/2} = (2.18 \pm 0.05) \cdot 10^{21} \text{ yr}$$

3.10. <sup>150</sup>Nd

The positive results were obtained in three independent experiments [39,61,82]. The most accurate value was obtained in Ref. [39]. This value is higher than in Ref. [61] (~  $3\sigma$  difference) and lower than in Ref. [82] (~  $2\sigma$  difference). Using Equations (2) and the three above-mentioned results, one obtains  $T_{1/2} = (8.4 \pm 0.5) \cdot 10^{18}$  yr. Taking into account that  $\chi^2/(N-1) > 1$  and S = 2.23 (see Equation (3)), we then obtain:

$$T_{1/2} = (8.4 \pm 1.1) \cdot 10^{18} \text{ yr}$$

It can be seen that due to the discrepancy between the  $T_{1/2}$  values, one has to increase the error in order to somehow agree on the experimental results. On the other hand, it is clear that the result of the NEMO-3 experiment is today the most accurate and reliable. This is confirmed by the fact that in the NEMO-3 experiment, seven different isotopes were investigated simultaneously. In addition to <sup>150</sup>Nd, <sup>48</sup>Ca, <sup>82</sup>Se, <sup>96</sup>Zr, <sup>100</sup>Mo, <sup>116</sup>Cd and <sup>130</sup>Te were also studied. For all these isotopes, the results are in good agreement with the results of other experiments. It is natural to assume that the result for <sup>150</sup>Nd is correct too. Therefore, I think that it is necessary to use this value as the most accurate at the moment:

$$T_{1/2} = 9.34^{+0.67}_{-0.64} \cdot 10^{18} \,\mathrm{yr}.$$

3.11.  ${}^{150}Nd - {}^{150}Sm (0^+_1; 740.4 \text{ Kev})$ 

There are two positive results for  $2\nu\beta\beta$  decay of <sup>150</sup>Nd to the 0<sup>+</sup><sub>1</sub> excited state of <sup>150</sup>Sm [83,84] (the preliminary result of Ref. [83] was published in Ref. [99]). These two results are in good agreement. The weighted average value is:

$$T_{1/2} = 1.2^{+0.3}_{-0.2} \cdot 10^{20} \text{ yr.}$$

Recently, the result of a new experiment was presented at MEDEX'19 [40] (see Table 1). I am not using this new result in my analysis because this is an ongoing experiment and the result is still preliminary and not yet published.

# $3.12.\ ^{238}U$

The two-neutrino decay of <sup>238</sup>U was measured in a single experiment using the radiochemical technique [85]:

$$T_{1/2} = (2.0 \pm 0.6) \cdot 10^{21} \text{ yr}.$$

It has to be stressed that for <sup>238</sup>U a "positive" result was obtained in only the experiment. Therefore, it is necessary to confirm this result in independent experiments (including direct measurements). Until these confirmations are received, one has to be very careful with this value.

## 3.13. <sup>130</sup>Ba (ECEC)

For <sup>130</sup>Ba, positive results were obtained using the geochemical technique only. In this type of measurement, one can not recognize the different modes. It is clear that exactly the ECEC(2 $\nu$ ) process was detected because other modes are strongly suppressed (see estimations in [91,100,101]). The first time the positive result for <sup>130</sup>Ba was mentioned was in Ref. [86], where experimental data of Ref. [102] were analyzed. In this paper, a positive result was obtained for one sample of barite ( $T_{1/2} = 2.1^{+3.0}_{-0.8} \cdot 10^{21}$  yr), but for a second sample only the limit was set ( $T_{1/2} > 4 \cdot 10^{21}$  yr). Later, more accurate half-life values, ( $2.2 \pm 0.5$ )  $\cdot 10^{21}$  yr [11] and ( $0.60 \pm 0.11$ )  $\cdot 10^{21}$  yr [87], were measured. One can see that the results are in strong disagreement. In [41], the data of [87] were analyzed and it was shown that subtraction of the contribution of cosmogenic <sup>130</sup>Xe removes the contradiction with the result of [11]. Finally, I recommend the following value from [11]:

$$T_{1/2} = (2.2 \pm 0.5) \cdot 10^{21} \text{ yr}$$

To obtain more reliable and precise half-life values, new measurements are needed (including direct experiments).

3.14.  $^{78}Kr(2K)$ 

The first indication of the observation of 2*K* capture in <sup>78</sup>Kr was announced in 2013 (the effect is  $\sim 2.5 \sigma$ ),  $T_{1/2} = [0.92^{+0.55}_{-0.26}(stat) \pm 0.13(syst)] \cdot 10^{22}$  years [103]. Then, the same data were analyzed more carefully and a new value was published ( $\sim 4 \sigma$ ), which turned out to be twice as much,

 $T_{1/2} = [1.9^{+1.3}_{-0.7}(stat) \pm 0.3(syst)] \cdot 10^{22}$  [13]. The analysis of the data is quite complicated and it is possible that the systematic error is much larger than the indicated 15%.

There is one more circumstance that makes me cautious about the result given in [13]. As can be seen from Table 3, in the case of <sup>78</sup>Kr, we are dealing with an anomalously large value of nuclear matrix element. This value is significantly larger than in the case of <sup>130</sup>Ba and <sup>124</sup>Xe (1.8 and 5.4 times, respectively) and exceeds all 13 NME values for  $2\nu\beta\beta$  decay (from 1.7 to 17.7 times). Here, it is necessary to take into account that, since the rate of the ECEC process is ~ 15–20% higher than the 2*K* capture, the NME for the ECEC process in <sup>78</sup>Kr is approximately 1.07–1.1 times greater than for the 2*K* capture. This circumstance only strengthens the contradiction. In principle, such a large NME is possible, but looks strange. In any case, confirmation of the result [13] in independent measurements is necessary. Until the confirmation, one has to be very careful with this result.

## 3.15. <sup>124</sup>Xe (2k)

To date, only one positive result has been published for 2*K* capture in <sup>124</sup>Xe [12]:  $T_{1/2} = [1.8 \pm 0.5(stat) \pm 0.13(syst)] \cdot 10^{22}$  yr. The significance of the effect is only 4.4  $\sigma$ . It should also be noted that a limit  $T_{1/2} > 2.1 \cdot 10^{22}$  yr was obtained in [104], which formally contradicts the result of [12]. Taking into account errors, there is no real contradiction here. However, it is clear that it is necessary to confirm the result of [12] in an independent experiment.

## 4. NME Values for Two-Neutrino Double Beta Decay

Obtained average and recommended half-life values are presented in Table 3 (2-nd column). Using these values, one can extract the experimental nuclear matrix elements through the relation [24]:

$$T_{1/2}^{-1} = G_{2\nu} \cdot g_A^4 \cdot (m_e c^2 \cdot M_{2\nu})^2, \tag{5}$$

where  $T_{1/2}$  is the half-life value in [yr],  $G_{2\nu}$  is the phase space factor in [yr<sup>-1</sup>],  $g_A$  is the dimensionless axial vector coupling constant and  $(m_ec^2 \cdot M_{2\nu})$  is the dimensionless nuclear matrix element. One has to remember that there are indications that in nuclear medium the  $g_A$  value is reduced in comparison with their free nucleon values (see Section 1). Expression (5) is valid for  $2\nu\beta\beta$  and ECEC( $2\nu$ ) processes.

Thereby, following Ref. [24], it is better to use the so-called effective NME,  $|M_{2\nu}^{eff}| = g_A^2 \cdot |(m_e c^2 \cdot M_{2\nu})|$ . This value has been calculated for all isotopes.

The obtained results are presented in Table 3 (3-rd and 4-th columns). When calculating, I used the  $G_{2\nu}$  values from Refs. [24,25] (see Table 4). For <sup>130</sup>Ba, <sup>78</sup>Kr and <sup>124</sup>Xe  $G_{2\nu}$  values for ECEC transition were taken from [25,105]. These calculations are most reliable and correct at this moment. The results of these calculations are in reasonable agreement (~1–7%) with three exceptions: for <sup>128</sup>Te (~20%), <sup>78</sup>Kr (~30%) and <sup>238</sup>U (factor ~ 7). For <sup>238</sup>U, two different values  $14.57 \cdot 10^{-21}$  yr<sup>-1</sup> [24] and  $98.51 \cdot 10^{-21}$  yr<sup>-1</sup> [25]) were produced. The situation with calculations for <sup>238</sup>U is clearly unsatisfactory and these calculations should be rechecked. For <sup>100</sup>Mo, <sup>100</sup>Mo-<sup>100</sup>Ru(0<sub>1</sub><sup>+</sup>) and <sup>116</sup>Cd, I used  $G_{2\nu}$  calculated in Ref. [24] for the SSD mechanism. The obtained values for  $|M_{2\nu}^{eff}|$  are given in Table 3 and these are the most correct values for these isotopes. So-called recommended values for  $|M_{2\nu}^{eff}|$  are given in Table 3 (5-th column) too. These values were obtained as an average of two values, given in columns 3 and 4. The recommended value error is chosen to cover all ranges of values from columns 3 and 4 (taking into account corresponding errors). For <sup>100</sup>Mo, <sup>100</sup>Mo-<sup>100</sup>Ru(0<sub>1</sub><sup>+</sup>) and <sup>116</sup>Cd, I recommended values obtained with  $G_{2\nu}$  for the SSD mechanism.

Therefore, for the majority of isotopes an accuracy for  $|M_{2\nu}^{eff}|$  is on the level ~1–10%. For <sup>78</sup>Kr, <sup>124</sup>Xe and <sup>130</sup>Ba, it is ~46%, ~22% and ~14%, respectively. This is mainly because of the not precise half-life values obtained for these isotopes. The most unsatisfactory situation is for <sup>238</sup>U (~70%). Main uncertainty in this case is connected with the accuracy of  $G_{2\nu}$ .

Isotope	$T_{1/2}(2\nu)$ , yr	$  M_{2\nu}^{eff}  $ ( $G_{2\nu}$ from [24])	$  M_{2\nu}^{eff}  $ (G <sub>2<math>\nu</math></sub> from [25])	Recommended Value
2νββ:				
<sup>48</sup> Ca	$5.3^{+1.2}_{-0.8}\cdot 10^{19}$	$0.0348\substack{+0.0030\\-0.0034}$	$0.0348\substack{+0.0030\\-0.0034}$	$0.035\pm0.003$
<sup>76</sup> Ge	$(1.88 \pm 0.08) \cdot 10^{21}$	$0.1051\substack{+0.0034\\-0.0023}$	$0.1074^{+0.0024}_{-0.0022}$	$0.106\pm0.004$
<sup>82</sup> Se	$0.87^{+0.02}_{-0.01}\cdot 10^{20}$	$0.0849^{+0.0005}$	$0.0855\substack{+0.0022\\-0.0010}$	$0.085\pm0.001$
<sup>96</sup> Zr	$(2.3 \pm 0.2) \cdot 10^{19}$	$0.0798^{+0.0010}_{-0.0037}$	$0.0804^{+0.0038}_{-0.0033}$	$0.080 \pm 0.004$
<sup>100</sup> Mo	$7.06^{+0.15}_{-0.13} \cdot 10^{18}$	$0.2071 \pm 0.0019$	$0.2096^{+0.0033}_{-0.0022}$	
1110	-0.13	0.2071 - 0.0022 0.1852 $+ 0.0017^{(a)}$	-0.0022	$0.185 \pm 0.002$
<sup>100</sup> Mo-	c <del>7</del> +0.5 1020	$\begin{array}{c} 0.1852\substack{+0.0017\ (a)\\-0.0019}\\ 0.1571\substack{+0.0048\\-0.0056}\end{array}$	0.1 < 10 + 0.0050	$0.185\pm0.002$
	$6.7^{+0.5}_{-0.4}\cdot 10^{20}$		$0.1619\substack{+0.0050\\-0.0058}$	
$^{100}$ Ru $(0^+_1)$	10	$0.1513^{+0.0047}_{-0.0053}{}^{(a)}_{-0.0053}$		$0.151\pm0.005$
<sup>116</sup> Cd	$(2.69\pm 0.09)\cdot 10^{19}$	$0.1160^{+0.0020}_{-0.0019}$	$0.1176\substack{+0.0020\\-0.0019}$	
		$0.1084^{+0.0024}_{-0.0019}$		$0.108\pm0.003$
<sup>128</sup> Te	$(2.25 \pm 0.09) \cdot 10^{24}$	$0.0406\substack{+0.0008\\-0.0008}$	$0.0454\substack{+0.0009\\-0.0009}$	$0.043\pm0.003$
<sup>130</sup> Te	$(7.91 \pm 0.21) \cdot 10^{20}$	$0.0288\substack{+0.0004\\-0.0004}$	$0.0297\substack{+0.0004\\-0.0004}$	$0.0293 \pm 0.0009$
<sup>136</sup> Xe	$(2.18 \pm 0.05) \cdot 10^{21}$	$0.0177^{+0.0002}_{-0.0002}$	$0.0184^{+0.0002}_{-0.0002}$	$0.0181 \pm 0.0006$
<sup>150</sup> Nd	$(9.34 \pm 0.65) \cdot 10^{18}$	$0.0543^{+0.0020}_{-0.0018}$	$0.0550\substack{+0.0020\\-0.0018}$	$0.055\pm0.003$
<sup>150</sup> Nd-	$1.2^{+0.3}_{-0.2} \cdot 10^{20}$	$0.0438\substack{+0.0018\\-0.0042}$	$0.0450\substack{+0.0018\\-0.0043}$	$0.044 \pm 0.005$
$^{150}$ Sm(0 <sup>+</sup> <sub>1</sub> )	-0.2	-0.0046	-0.0048	
<sup>238</sup> U	$(2.0\pm0.6)\cdot10^{21}$	$0.1853\substack{+0.0361\\-0.0227}$	$0.0713\substack{+0.0139\\-0.0088}$	$0.13\substack{+0.09 \\ -0.07}$
ECEC( $2\nu$ ):			0.0000	0.07
<sup>78</sup> Kr <sup>(b)</sup>	$1.9^{+1.3}_{-0.8} \cdot 10^{22}$	$0.2882^{+0.0829}_{-0.0706}$ [105]	$0.3583^{+0.1126}_{-0.0822}$	$0.32\substack{+0.15 \\ -0.11}$
$^{124}$ Xe $^{(b)}$	$(1.8 \pm 0.5) \cdot 10^{22}$	$0.0568^{+0.0101}_{-0.0650}$ [105]	$0.0607^{+0.0107}_{-0.0070}$	$0.059\substack{+0.013\\-0.009}$
<sup>130</sup> Ba	$(2.2 \pm 0.5) \cdot 10^{21}$	$0.1741^{+0.0239}_{-0.0170}$ [105]	$0.1754^{+0.0241}_{-0.0171}$	$0.175_{-0.017}^{+0.024}$

**Table 3.** Half-life and effective nuclear matrix element values for  $2\nu\beta\beta$  decay (see Section 4).

 $^{(a)}$  Obtained using the SSD model.  $^{(b)}$  Value for 2K capture. For the ECEC process, the half-life value will be approximately 15–20% less, and the NME value approximately 7–10% higher.

Recently, in Ref. [16], an improved formalism of the  $2\nu\beta\beta$  decay rate was presented, which takes into account the dependence of energy denominators on lepton energies via the Taylor expansion. As a result, the formula for the half-life starts to be more complicated and contains several different matrix elements and different phase space volumes. That is, a new approach to processing the results will be required. To do this, some parameters of this approach have to be established from experiment and calculated reliably, e.g., within the interacting shell model (see discussion in [16]). Nevertheless, the results shown in Table 3 retain their significance since it was demonstrated in [16] that additional terms contribute ~3% to ~25% to the total decay rate. This means that if we consider expression (5) as the first term of the expansion in the approach [16], then we can conclude that the values of  $|M_{2\nu}^{eff}|$ obtained in this work give a good estimate for  $g_A^2 \cdot |M_{GT-1}^{2\nu}|$  (see Formula (19) in [16]). The values given in Table 3 overestimate  $g_A^2 \cdot |M_{GT-1}^{2\nu}|$  values by ~1.5–12% only, which is comparable to the accuracy of determining  $|M_{2\nu}^{eff}|$ . An exception is the situation with the results for <sup>100</sup>Mo and <sup>116</sup>Cd obtained using phase space volumes calculated within the SSD. In this case, the most accurate NME estimate was obtained since the exact value of the energy of the lowest 1<sup>+</sup> intermediate state was used in the calculations of the phase space volume.

Isotope	$G_{2\nu}(10^{-21}{ m yr}^{-1})$ [24]	$G_{2\nu}(10^{-21}{ m yr}^{-1})$ [25]
2νββ:		
<sup>48</sup> Ca	15,550	15,536
<sup>76</sup> Ge	48.17	46.47
<sup>82</sup> Se	1596	1573
<sup>96</sup> Zr	6816	6744
<sup>100</sup> Mo	3308	3231
	4134 <sup>(a)</sup>	
$^{100}$ Mo- $^{100}$ Ru $(0^+_1)$	60.55	57.08
	$65.18^{(a)}$	
<sup>116</sup> Cd	2764	2688
	3176 <sup>(a)</sup>	
<sup>128</sup> Te	0.2688	0.2149
<sup>130</sup> Te	1529	1442
<sup>136</sup> Xe	1433	1332
<sup>150</sup> Nd	36,430	35,397
$^{150}$ Nd- $^{150}$ Sm(0 <sup>+</sup> <sub>1</sub> )	4329	4116
<sup>238</sup> U	14.57	98.51
ECEC( $2\nu$ ):		
<sup>78</sup> Kr	0.660 [105]	0.410
<sup>124</sup> Xe	17.200 [105]	15.096
<sup>130</sup> Ba	15.000 [105]	14.773

Table 4. Phase-space factors from Refs. [24,25,105].

<sup>(*a*)</sup> Obtained using SSD model.

#### 5. Conclusions

Thus, the all positive results for  $2\nu\beta\beta$  decay obtained by August 2020 have been analyzed. As a result, the average values of the half-life were obtained for all considered isotopes. For <sup>128</sup>Te, <sup>150</sup>Nd and <sup>130</sup>Ba, so-called recommended values have also been proposed. Using these obtained average/recommended half-life values, the  $|M_{2\nu}^{eff}|$  values for all considered nuclei were determined. Finally, previous results from Ref. [15] were successfully updated. A summary is shown in Table 3. I recommend using these values as the most correct and reliable currently. If we look at the dynamics of the average values since 2001, we can see that these values were constantly refined over time and did not deviate by more than 1–2  $\sigma$  from the initial value. An exception is the situation with <sup>76</sup>Ge. Here, the average value has steadily increased with time (from  $1.42^{+0.09}_{-0.07} \cdot 10^{21}$  yr in 2001 to  $(1.88 \pm 0.08) \cdot 10^{21}$  yr in 2020). This is due to the low quality of the results obtained in the 1990s. In the latest analysis, the results obtained after 2000 have been used.

At present,  $2\nu\beta\beta$  decay was recorded in 11 nuclei, and ECEC capture in 3 nuclei (with some doubts for <sup>78</sup>Kr). The accuracy of determining the half-life for most nuclei lies in the range 2–10%. It is expected that in the next few years new results will be obtained for <sup>76</sup>Ge (Majorana), <sup>100</sup>Mo (CUPID-Mo, AMORE, CROSS), <sup>116</sup>Cd (CROSS), <sup>130</sup>Te (CROSS, SNO+), <sup>136</sup>Xe (NEXT-100) and <sup>124</sup>Xe (NEXT-100, LUX-ZEPLIN). The final result will be obtained in an experiment to search for the  $2\nu\beta\beta$  decay of <sup>150</sup>Nd to the first excited 0<sup>+</sup> level of <sup>150</sup>Sm (see [40]). Let us emphasize here the importance of experiments using low-temperature bolometers. In experiments with such detectors, the measurement accuracy of the half-life can reach 1–2%. At present, such experiments are possible for <sup>82</sup>Se, <sup>100</sup>Mo, <sup>116</sup>Cd and <sup>130</sup>Te. Apparently, in the future, such measurements will be implemented for <sup>48</sup>Ca as well. I hope that in the future 2 $\beta$  processes will also be found in other nuclei. The search for  $2\nu\beta\beta$  processes in <sup>124</sup>Sn, <sup>110</sup>Pd, <sup>160</sup>Gd and the search for ECEC(2 $\nu$ ) processes in <sup>96</sup>Ru, <sup>106</sup>Cd and <sup>136</sup>Ce seem promising. As for the  $2\nu\beta\beta$  transitions to the excited states of the daughter nucleus, it seems possible to register a transition to the 0<sup>+</sup> excited level in measurements with <sup>96</sup>Zr and <sup>82</sup>Se in the near future.

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