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Special introductory note:

To see how the quantization in radioactive decay measurements was first discerned, please read the paper "Systematic Fractional Relationships in Radioactive Decay Measurements".

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SYSTEMATIC FRACTIONAL RELATIONSHIPS IN RADIOACTIVE DECAY MEASUREMENTS 28/05/11

(Quantization in Radioactive Decay Measurements). Refreshed 03/12/13

TABLE 1a The 2.41 group of radioactive nuclides.

Note the decay constant of 2.41

Wuclide)	Half Time:	5.005	tional tionship	Dec: Cons	ay stant
Light !	86.4	2/3 x	129.6		x 2.41
?	43.2	1/2 x	86.4	0	x 2.41
7	28.8	2/3 x	43.2		x 2,41
Nd 144	21.6	3/4 x	1000	٥	x 2,41
Th 232	14.4	2/3 x	21.6	125	x 2,41
U 235	7.2	1/2 x	14.4		x 2,41
Rb 87	4.8	·2/3 x	7.2		x 2.41
Lu 176	3.6	3/4 x	4.8	8	x 2.41
?	(2.4)	2/3 x	3.6	1.2	x 2.41
K 40	1.2	1/2 x	2.4.	2.4	x 2.41
Sm 148	.8	. 2/3 x	1.2	. 3.6	x 2.41
Pt 190	.6	3/4 x	.8	4.8	x 2.41
Re 187	.4	2/3 x	.6	7.2	x 2.41
Hf174/Te	130 .2	1/2 x	.4	14.4	x 2.41
?	-13	2/3 x	.2		x 2.41
3.	.1	3/4 x	.13		x 2.41

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TABLE 1b The 1.5 group of radioactive muclides. Note the decay constant of 1.5

		.1 2.00	
Nuclide	Half Time	Fractional Half Fine	Decay Constant
T 238	4.6	$\frac{10 \times 1}{9 \times 2.41}$	1.5
In 115	5.1	10 x 10 x 1 9 9 2.41	9 x 1.5
Od. 113.	9	9	$\frac{10 \times 10 \times 1.5 \times 1}{9} \times \frac{10}{9} \times 1.5 \times \frac{1}{2.41}$
Se 82 La 138 Sm 147	1.11	. 10 9	1.5 x 1 2.41
Gd 152			

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NOTES

- 1. 'Powers of ten' excluded. These may be found from the standard half life listings.
- 2. Becay constant = .693 / half life
 Half life = .693 / decay constant
- 3. R 40. Published decay constant = 5.8

 Half life = 1.2

 U 238. Published decay constant = 1.5

 Half life = 4.6
- 4. 2.41 approximate. Use 2.406(25)

given by ods of A

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constant, in exists, in exists, in autity, in T_B , etween a lucts in a uranium t equilib- 4.5×10^9 ; radioacween uram shows the to be ity of the

5) shows ed unity; illibrium.

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in the active n at $t_m = few$ hunts parent, 07/(307-40)Ba.

'TABLE' A. Parent radioactive nuclides found in nature

Nuclide		9			K DESCRIPTION FILE IN SECTION	
Atamic númber, Z	Mass number, A	Percent abundance in nature	Half-period, years	Radioactive transitions observed*	Disintegration energy, MeV*	
19.K	40	0.0117	1.3 × 10 ⁹	β-, β*, EC	β 1.3 - EC 1.5	
34 Se	82	9.19	1.1×10^{20}	β-β-	3.0	
37 Rb	87	27.83	4.8×10^{10}	β-	0.3	
48 Cd	113	12.2	9×10^{15}	β^-	0.3	
49 In	115	95.77	5.1×10^{14}	<i>β</i> -	0.5	
52 Te	130	34.49	2×10^{21}	Growth of 130 Xe1	1.6	
57 La	138	0.089	1.1×10^{11}	β^- , E \tilde{G}	β^- 1.0 – EC 1.75	
60 Nd	144	23.8	2.1×10^{15}	α	1.9	
62 Sm	147	15.07	1.1×10^{11}	a	2.3	
62 Sm	148	11.3	8×10^{15}	α	1.99	
64 Gd	152	0.20	1.1×10^{14}	O:	2.2	
71 Lu	176	2.6	3.6×10^{10}	β^-, γ	0.6	
72 Hf	174	0.16	2×10^{15}	α	2.5	
75 Re	187	62.6	4×10^{10}	β^{-}	0.003	
78 Pt	190	0.013	6×10^{11}	α	3.24	
90 Th	232	100	1.4×10^{10}	O.	4.08	
92 U	235	0.715	7.0×10^{8}	a, SF	α 4.68	
92 U	238	99.28	4.5×10^{9}	a, SF	cx 4.27	

EC = electron capture. The EC energy is between ground states, but, in ¹⁹⁸La decay, 1.44 MeV of the EC energy goes to a gamma ray that feeds the ground state.

Indirect evidence for \$\beta^-\beta^-\decay.

Radioactivity in the Earth. A number of isotopes of elements found in the Earth are radioactive (Table 2). All known or theoretically predicted isotopes of elements above bismuth are radioactive. Because the Earth is composed of atoms which were believed to have been created more than 3 × 109 years ago, the naturally occurring parent radioactive isotopes are those which have such long half-periods that detectable residual activity is still observable today. As a general rule, one can detect the presence of a radioactive substance for about 10 half-lives. Therefore activities with $T\lesssim 0.3$ × 109 years should not be found in the Earth. For example, present-day uranium is an isotopic mixture containing 99.3% ²³⁸U, whose half-period is 4.5 x 109 years, and only 0.7% of the shorter-lived uranium isotope 235 U, whose half-period is 0.7×10^9 years, whereas these isotopes presumably were produced in roughly equal amounts in the Earth a few billion years ago. Geophysical evidence indicates that originally some 236U was present also, but none is found in nature now as expected with its half-period of 0.02×10^9 years. The elements technetium (Z = (3) and promethium (Z = 61) are not found in the l'arth's crust because all their isotopes are radioactive with much shorter half-periods (their longest-lived are $T=2.6 \times 10^6$ years for ⁹⁷Tc and T=17.7 years for 145Pm).

Uranium-238 decays through a long series of 14 radioactive decay products before ending as a stable isotope of lead, ²⁰⁶Pb. Some of these members of the ²³⁸U decay chain have very short half-periods, so their existence in nature is entirely dependent on the presence of their long-lived parent, and thus is a senealogical accident. For example, radium occurs in nature only in the minerals of its parent, uranium. The decay series of ²³⁵U supports 14, and the decay

series of 232 Th supports 10, short-lived radioactive substances found in nature.

A few of the common elements contain long-lived, naturally radioactive isotopes. For example, all terrestrial potassium contains 0.012% of the radioactive isotope $^{40}\mathrm{K}$, which has a half-period of 1.3 \times 10^9 years, and emits negatron or positron beta particles (plus decay via electron capture) and gamma rays in a dual decay to stable $^{40}\mathrm{Ca}$ and $^{40}\mathrm{Ar}$. This isotope is the principal source of radioactivity in a normal human being; each human contains about 0.1 microcurie (3.7 \times 10^3 becquerels) of the radioactive potassium isotope $^{40}\mathrm{K}$.

Geological age measurements are based on the accumulation of decay products of long-lived isotopes, especially in the cases of ⁴⁰K, ⁸⁷Rb, ²³²Th, ²³⁵U, and ²³⁸U. *See* GEOCHRONOMETRY.

Laboratory-produced radioactive nuclei. With particle accelerators and nuclear reactors, the order of 2000 radioactive isotopes not found in detectable quantities in the Earth's crust have been produced in the laboratory since 1935, including those of 26 new chemical elements up to element 118 (as of 2004), with element 117 not known. Earlier titles of induced or artificial radioactivities for these isotopes are misnomers. Many of these now have been identified in meteorites and in stars, and others are produced in the atmosphere by cosmic rays. There are over 5000 isotopes theoretically predicted to exist. As one approaches the place where a proton or neutron is no longer bound in a nucleus of an element (the limits of the existence of that element), the halfperiods become extremely short. See TRANSURANIUM

For example, carbon-14 is a negatron beta-particle emitter, with a half-period of about 5600 years, which can be produced in the laboratory as the

12 14

e product, vhose

Quantization in the radioactive decay rates of the members of the three naturally occurring radioactive transformation series. Compiled 13/1/2015

(Compa	re Table	1a).	•			Compile	ed 13/1/2015
Parent Nuclide	Transformation Series Member	Quantized 差 Time Position	Quantized Decay Rate x 2.406(25)	Ealf Time	LANGE TO COLL TO	Fractional Relationships of New 2 Times e.g.	57.6/86.4 = 2/3 43.2/57.6 = 3/4 Comments
Light	Ra228 Po218 Po215	172.8 129.6 86.4 (57.6)	.033 .05 " .066	- 5.8 5.7x 5.7x	10 ⁻⁶ 10 ⁻¹¹		nt Science News ollo.org.nz
Nd144 Th232 U235	Ac227 Pb210 Th228 Bi215	21.6 (19.2) 14.4 7.2	.133 " .15 .2	100	x 10 ⁻⁵		of Auckland, N.Z.
Rb87	At218 Po216 Pa234 m Rn220 Ra226	4.8 3.6 2.4 (1.8) (1.6)	.6 .8 1.2 1.6	2.28	x 10 ⁻⁸ x 10 ⁻⁹ x 10 ⁻⁶ x 10 ⁻⁶	3/4, 2/3 2/3, 3/4	position?
K40 Sm148	T1206	1.2 11 .8	2.4 11 3.6	.00. .000 80,000 7.99 5.89	× 10 ⁻⁶ × 10 ⁻⁶	1.1	on Table 1b?
Re187 Hf174 Te130	Fr223 Bi211 Th231 Ra223	.4 (.3)	7.2 9.6 n	4.18 4.18 .002 .030			position? position?
	Bi210 Ra224 Rn222 Rn218 Po212 Tl207 Pa234 Th234	.133 .1 " " " (.075)	21.6 28.8 "" "" 38.4 43.2	9.51 9.13 .000	01 04 × 10 10 × 10 15 × 10 6 0764		.1 on Table 1b? /10 on Table 1b?
e e	Ac228 Pb211 Th227 Pb214 Po214 Bi214 Po210 Pa231 U234	(.05) (.05) (.0375) (.033 (.025)	57.6 11 78.6 11 86.4 115.2	6.84 .052 5.13 5.07 3.80 .378	x 10 ⁻¹ 2 x 10 ⁻⁵ x 10 ⁻⁵	88 . 23 26 88	5.1 on Table 1b? " " " " " " " " " " " " 9: (.0375/.05)
42	T1210 At219 Rn219	.022 .0166 (.0125)	129.6 172.8 230.4	2.47 1.71 1.27		Or, 172.8	position? 29.6 position? ble 1b

TABLE B. Names, symbols, and radioactive properties of members of the three naturally occurring radioactive transformation series.

Early name [†]	Early symbol [†]	Atomic number	Mass number	Isotopic symbol	Half- period	Type o decay
		Uṛa	nium (4n + 2) serie			
Uranium I	UI	92	238	238U	$4.5 \times 10^{9} \text{ y}$	a, SF
Uranium X ₁	UX ₁	90	234	234Th	24 d	B
Ufanium X ₂	UX ₂	91	234	234mpa	1.2 m	17,,3-
Uranium Z	UZ	91	234	234 Pa	6.7 h	<i>8</i> -
Uranium II	Uil	92	234	234[]	$2.5 \times 10^5 \text{ V}$	α
ionium	lo	90	230	286 Th	$8 \times 10^{4} \text{y}$	O!
Radium	Ra	88	226	²²⁶ Ra	1600 y	α
Radon	Rn	86	222	222 Rn	3.8 d	· or
Radium A	RaA	84	218	218pg	3.0 m	α, β^-
Astatine	At	85	218	218 At	1.5 \$	β^{-} , α
Radon	Pin	86	218	218Rn	3,5 ms	ά
Radium B	RaB	82	214	214Pb	27 m	\mathcal{B}^-
Radium C	RaC	83	214	²¹⁴ Bi	20 m	β-, α
Radium C'	RaC'	84	214	214Po	$1.6 \times 10^{-4} \text{ s}$	CΣ
Radium C"	RaC"	81	210	21077	1.3 m	B-
Radjum D	RaD	82	210	210Pb	22 y	<i>Ġ</i> ~
Radium E	RaE	83	210	210Bi	5.0 d	β-, ο
Radium F	RaF	84	210	210 Po	138 d	O
Thallium	TI	81	206	206∏	4.2 m	β^-
Radium G	RaG	82	206	208 Pb	Stable	Stabl
Total Control Control			horium (4n) series		3.73.77	,
TL 2	Th	90	232	²³² Th	$1.4 \times 10^{10} \text{y}$	W 827
Thorium	MsTh₁	90 88	232	²²⁸ Ra	5.8 y	α 3-
Mesothorium ₁	12.5		228	228 AG	6.1 h	β-
Mesothorium ₂	MsTh ₂	89		228Th	1.9 y	- 50
Hadiothorium	RdTh	90	228	²²⁴ Ra	3.7 d	O
Thorlum X	ThX	88	224	220 Rn		CY
Thoron	Tn	86	220	216Po	56 s	CK
Thorium A	ThA	84	216	212Pb	0.15 s	α
Thorium B	ThB	82	212	212Bi	10.6 h	β-
Thorium G	ThC	83	212		1.0 h	β , α
Thorium G'	ThO'	84	212	²¹² Po ²⁰⁸ TJ	3 × 10 ⁻⁷ s	a.
Thorium C*	ThC"	81	208		3.1 m	β-
Thorium D	ThD	82	208	²⁰⁸ Pb	Stable	Stabl
		Acti	nium (4n + 3) serie			
Actinouranium	AcU	92	235	235U	$7.0 \times 10^{8} \mathrm{y}$	a, SF
Uranium Y	UY ·	90	231	231 Th	26 h	β^{-}
Protactinium	Pa	91	231	231 Pa	$3.3 \times 10^4 \text{ y}$	CE.
Actinium	Ac	89	227	²²⁷ Ac	22 y	β^- , α
Radioactinium	RdAc	90	227	²²⁷ Th	19 d	(X
Actinium K	AcK	87	223	223 Fr	22 m	β^- , o
Actinium X	AcX	88	223	²²³ Ra	11 d	α
Astatine	At	85	219	²¹⁹ At	0.9 m	α, β^-
Actinon	An	86	219	219 Rn	4.0 s	α
Bismuth	Bi	83	215	²¹⁵ Bi	7.6 m	β^{-}
Actinium A	AcA	84	215	215Po	$1.8 \times 10^{-3} \text{ s}$	ćv.
Actinium B	AcB	82	211	211 Pb	36 m	β-
Actinium C	AcC	83	211	211Bi	2.2 m	α, β
Actinium G'	AcG'	84	211	211 Po	0.5 s	α.
Actinium C"	AcC"	81	207	207TI	4,8 m	ß
Actinium D	AcD	82	207	207 Pb	Stable	Stabl

Radon-223 has been shown to have a very weak radioactive ¹⁴C decay branch to ²⁰⁹Pb. Several of the isotopes in these chains, such as ^{234,235,236}U, ²³Pa, and others are predicted to have such very weak, heavy cluster decay branches.

These were the names and symbols used before the different isotopes of these elements were known.

Transformation series are now known for every element in the periodic table except hydrogen. Chains of neutron-rich isotopes have been produced and studied among the products of nuclear fission. Heavy-ion-induced reactions and high-flux reactors have been used to extend knowledge of the elements beyond uranium. The elements from number 93 (neptunium) to 118 (as yet unnamed except for element 117), which have so far not been found on flarth, were made in the laboratory. Both proton- and heavy-ion-induced reactions have extended knowl-

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edge of chains and neutron-deficient isotopes of the stable elements.

Alpha-Particle Decay

Alpha-particle decay is that type of radioactivity in which the parent nucleus expels an alpha particle (a helium nucleus). The alpha particle is emitted with a speed of the order of 1 to 2×10^7 m/s (10^4 mi/s), that is, about 1/20 of the velocity of light.

In the simplest case of alpha decay, every alpha particle would be emitted with exactly the same