

NOTIZEN

Comment on the Age of the Elements

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REYNOLDS¹ has recently discovered that the isotopic composition of xenon in the chondrite Richardton is different from that of atmospheric xenon. The most prominent deviation from terrestrial xenon is an excess of Xe¹²⁹ of about 1.3 · 10⁻¹⁰ ccSTP/g, produced from the decay of I¹²⁹ (half-life 17 · 10⁶ years). By assuming that this Xe was produced inside the Richardton meteorite, REYNOLDS deduced that the maximum time which elapsed between the synthesis of the elements and the formation of Richardton was 3.5 · 10⁸ years.

In this note we wish to stress some arguments which would favour an alternative conclusion with respect to the time of formation of the elements. All the xenon, including the excess Xe¹²⁹, could be trapped primeval gas from an atmosphere in which the decay of the I¹²⁹ contributed relatively more Xe¹²⁹ than in our atmosphere.

Table 1 shows the light rare gases in some stone meteorites. In chondrites all of them, except A³⁶, can be explained by radioactive decay or spallation^{2, 3}. The reaction Cl³⁵(n, γ)Cl³⁶ → A³⁶ might provide for some of the excess A³⁶. To explain the total A³⁶ excess in some of the meteorites (Bjurböle, Richardton), however, the required thermal neutron fluxes would have to be high and could hardly be provided by cosmic ray interactions. Also, the low He³/A³⁸ ratio of Bjurböle is not explained, because the reaction Cl³⁷(n, γ)Cl³⁸ → A³⁸ has a much too low cross-section.

A possible explanation would be that small amounts of primeval argon were trapped in the crystals during the formation of the meteorites. In fact, GERLING and

LEVSKII⁵ found in the Pesyanoe achondrite large amounts of primeval helium, neon, and argon and the excessive amount of A³⁶ in Richardton is only about 1% of the primeval A³⁶ in Pesyanoe. The content of rare gases in Pesyanoe shows that this meteorite was formed in an atmosphere in which the rare gases had a high partial pressure. Furthermore, from the abundances of helium and neon relative to A³⁶, it is evident that greater fractionation has occurred in the terrestrial atmosphere than in the "Pesyanoe atmosphere" (Table 2). If the excess A³⁶ is primeval, the absence of measurable amounts of primeval neon in Richardton would require a still greater fractionation, in agreement with a relatively high ratio Xe/A³⁶ in Richardton.

The evidence available so far seems to indicate that the occurrence of trapped primeval gas has to be taken into account. It also seems quite reasonable that primeval xenon may have different Xe¹²⁹ contents in different environments. Therefore, we think the assumption, that the excess Xe¹²⁹ was formed inside the meteorite, cannot be made with certainty at present, and the time between nucleogenesis and meteorite formation may have been longer than 0.35 · 10⁹ years.

	He ⁴	Ne ²⁰	A ³⁶	Kr	Xe
Richardton	—	<.2	1	—	≤.05
Pesyanoe	4000	14	1	—	—
Terrestrial atmosphere	—	.52	1	.032	.0025
Cosmic abundance	2.4 × 10 ⁴	62	1	.00041	.000032

Table 2. Relative Abundance of "Primeval Gases". Primeval gas contents calculated from Table 1 and from REYNOLDS¹. Composition of the terrestrial atmosphere from RANKAMA and SAHAMA⁶, cosmic abundances from SUESS and UREY⁷.

	A ³⁶ × 10 ⁻⁸ ccSTP/g	A ³⁶ /A ³⁸	Ne ²⁰ × 10 ⁻⁸ ccSTP/g	Ne ²⁰ /Ne ²¹	He ⁴ × 10 ⁻⁶ ccSTP/g	He ³ /A ³⁸
Holbrook (Ch)	1.26	1.18	6.45	.965	18.3	26
St. Michel (Ch)	1.94	1.83	7.95	.97	6.25	30
Richardton (Ch)	2.83	2.44	9.10	.96	15.1	29
Bjurböle (Ch)	4.62	3.80	4.40	.99	16.7	14
Furnas County (A)	1.83	.93	56.5	.965	14.1	
Pasamonte (A)	.66	.72	—	—	61	
Pesyanoe (A)	165	5	2200	110	6500	
Terrestrial atmosphere	—	5.35	—	350	—	

Table 1. Light rare gases in some stone meteorites. Ch = chondrite; A = achondrite. Argon data from GEISS and HESS²; neon from P. and A. EBERHARDT⁴; helium from EBERHARDT and HESS³. All data of Pesyanoe from GERLING and LEVSKII⁵.

¹ J. H. REYNOLDS, Phys. Rev. (Letters) 4, 8 [1960].

² J. GEISS and D. C. HESS, Astrophys. J. 127, 224 [1958].

³ P. EBERHARDT and D. C. HESS, Astrophys. J. 131, 38 [1960].

⁴ P. and A. EBERHARDT, to be published in Z. Naturforschg.

⁵ E. K. GERLING and L. K. LEVSKII, Dokl. Akad. Nauk, SSSR 110, 750 [1956].

⁶ K. RANKAMA and Th. G. SAHAMA, Geochemistry, The University of Chicago Press 1950.

⁷ H. E. SUESS and H. C. UREY, Rev. Mod. Phys. 28, 53 [1956].



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