

of the very strong short-lived cobalt and manganese activities in these same iron samples. A particularly rigorous chemical separation, designed to remove all elements other than iron, was made on a more recently activated specimen; this precipitate, now 10 months old, has already departed from the 47-day period and is now going into one much longer.

These facts assure us that  $\text{Fe}^{55}$  has been formed through  $\text{Fe}^{54}(d, p)\text{Fe}^{55}$  with the activity probably leading to stable  $\text{Mn}^{55}$  either by positron emission or by  $K$ -electron capture. An absorption curve on one of the old samples (unfortunately very weak) shows that the radiation has a soft component with half-thickness 3 to 5  $\text{mg}/\text{cm}^2$  of aluminum. This suggests the existence of an internally converted gamma-ray.

It is too early to give a reliable value for the half-life of this activity. The observed decay curves, which are still going through the transition interval from the 47-day period to the long period, set a lower limit of 1 year for the half-life. A comparison of the intensities of the 47 day and the longer activity (assuming equal cross sections for their production and equal ionizing powers for their radiations, and allowing for the abundances of the primary isotopes involved) predicts that the half-life will exceed 10 years.

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<sup>1</sup> J. J. Livingood and G. T. Seaborg, *Phys. Rev.* **54**, 51 (1938).

<sup>2</sup> A. O. Nier, *Bull. Am. Phys. Soc.* **14**, 40 (1939).

<sup>3</sup> J. J. Livingood and G. T. Seaborg, *Phys. Rev.* **53**, 847 (1938).

<sup>4</sup> J. J. Livingood and G. T. Seaborg, *Phys. Rev.* **54**, 391 (1938).

#### Stability of Uranium and Thorium for Natural Fission

The question of the occurrence of possible natural fission processes among the heavy nuclei is one of considerable importance to both nuclear physics and geophysics. The characteristics of the neutron-induced splitting of the uranium and thorium nuclei which have recently been studied so intensively by many workers seem to indicate that a naturally occurring explosion process probably would form radioactive isotopes of elements of medium atomic number with convenient half-lives. Further, the emission of neutrons by uranium in the artificial process suggests that neutron emission might accompany the natural process. Finally, natural fission might be detected by disparities between the results of the various calculations of the age of the oldest rocks based on the observed abundances of the elements and the intensities of the ordinary radioactive emissions from uranium and thorium.

It is the purpose of this letter to point out how well evidence on these three points excludes natural fission. In all of the numerous experiments which have been done on the neutron-produced activities of uranium and thorium control separations must have been made without neutron

irradiation to exclude the possibility of contamination by natural activities. In order to have definite data for a typical calculation for this type of evidence the experiments<sup>1, 2</sup> on extraction of radioiodine from irradiated uranium were repeated with 0.90 mole of uranyl nitrate which had not been disturbed chemically for at least five years. The iodine, extracted by adding a little solid iodine to a slightly acidified aqueous solution of the uranyl nitrate, shaking with  $\text{CCl}_4$ , and precipitating  $\text{AgI}$  in a thin layer was counted in a screen wall counter so no beta-radiation harder than 20,000 electron volts would have been missed. The result was zero within 20 counts per minute. If it is assumed that each ten explosions produced at least one radioactive iodine on the average this result requires that the half-life for natural fission be at least as long as  $10^{14}$  years. Similar results must apply for the other elements which have been separated, so we can conclude that natural fission to produce any of the radioactive elements found in the artificial process must correspond to a half-life of the order of or greater than  $10^{14}$  years. A similar test was performed on thorium with the same results.

In order to investigate possible natural neutron emission from uranium and thorium a  $\text{BF}_3$ -filled counter was surrounded with paraffin and standardized with 200 mg of  $\text{RaBr}_2$  mixed with Be powder. Seven and one-half moles of uranium salts were found to give less than two counts per minute whereas the Ra-Be source in the same position gave 4600 counts per minute. If it is assumed that the Ra-Be source gives 17,000 neutrons per second per millicurie,<sup>3</sup> the half-life for fission must be at least as large as  $10^{14}$  years if at least one neutron were emitted per explosion. Similar measurements on a smaller amount of thorium gave a lower limit of  $5 \times 10^{13}$  years.

The agreement between the uranium and thorium age determinations on old rocks and the agreement of these with other estimates of the age of the oldest strata, e.g., the  $\text{Sr}^{87} : \text{Rb}^{87}$  ratio found by Hahn, Strassmann and Walling<sup>4</sup> in lepidolite from the Canadian shield indicate that the age of the earth is at least  $2 \times 10^9$  years and that no fission process with a half-life less than  $5 \times 10^9$  years occurs, unless uranium and thorium have been or are now being generated by some terrestrial process. In the absence of any evidence for such a generation we must conclude that the half-life for any possible fission exceeds  $5 \times 10^9$  years.

Consideration of all these points leads to the conclusion that no fission similar to those induced by neutron absorption can occur naturally for either uranium or thorium at a rate greater than that corresponding to a half-life of  $10^{14}$  years and that no natural fission of any sort has a life less than  $5 \times 10^9$  years.

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<sup>1</sup> P. Abelson, *Phys. Rev.* **55**, 670 (1939).

<sup>2</sup> R. W. Dodson and R. D. Fowler, *Phys. Rev.* **55**, 880 (1939).

<sup>3</sup> E. T. Booth, J. R. Dunning and F. G. Slack, *Phys. Rev.* **55**, 876 (1939). Their figure of 15,000 for radon has been increased for the extra alpha-particle from radium.

<sup>4</sup> O. Hahn, F. Strassmann and E. Walling, *Naturwiss.* **25**, 189 (1937). F. Strassmann and E. Walling, *Berichte* **71**, 1 (1938).