Absorption Curve of 31-Second ₈O¹⁹ Beta-Rays and Cross Section for Production by **Thermal Neutrons***

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THE absorption of 31-second 8019 beta-rays has been measured by irradiating ordinary distilled water in the Argonne pile. The beta-rays were counted on a Duraluminum-walled Geiger counter. Aluminum absorbers up to 0.875 gram per cm² were used, and one point was taken with a 1.84 gram per cm² Pb absorber. (See the attached graph, Fig. 1.) From the absorption curve the range can be estimated as ~ 1.4 grams per cm² Al which corresponds to 3 Mev beta-rays. A further check on the energy is obtained by comparing the mass absorption coefficient of the 8O19 beta-rays with that from 13Al28 and 19K42 beta-rays which we measured on the same geometry. For 31 sec. $_{8}O^{19}$ beta-rays, we got u = 2.56 cm² per gram Al. For 2.4 min $_{13}$ Al²⁸ beta-rays, we got u = 2.5 cm² per gram Al, and a cloud-chamber measurement¹ of the energy of these betarays gave 3.4 Mev. For 12.4 hr. 19K42 beta-rays, we got u = 2.56 cm² per gram Al, and a cloud-chamber measurement² of the energy of these beta-rays gave 3.5 Mev.

From the range and absorption coefficient data, it would seem reasonable to conclude that the energy of the ₈O¹⁹ beta-rays is about 3.3 Mev. Gamma-rays also accompany the 31-sec. activity, as shown by the point on the absorption curve taken with the Pb absorber.

In addition to the 31-sec. beta-rays, we observed a half-life of ~ 4 minutes, probably caused by an impurity in the water. The saturation value of the \sim 4-minute activity was \sim 25 percent of the 31-sec. activity.

John Marshall³ has measured the thermal neutron activation cross section⁴ of ₈O¹⁸ assuming that ₈O¹⁹ betarays had the same absorption coefficient as UX_{II} beta-rays. On the same geometry that we used to measure the absorption coefficient of 8019 beta-rays, we obtained $\mu = 4.88$ cm² per gram for UX_{II} beta-rays. Hence, Marshall's value of 3.5×10^{-4} barn⁵ for the thermal neutron activation cross section of ₈O¹⁸ is high because he assumed $_{8}O^{19}$ beta-rays to be less penetrating than they actually are. A correction was made as follows: Counter wall used



FIG. 1. Absorption of B-rays from O19.

in Marshall's measurements was 0.050 gram per cm² Al, thickness of water layer used was 3 mm or 0.3 gram per cm². One-half of the water layer was the effective selfabsorption thickness. Hence, the beta-ray absorption correction factor

 $\exp^{-(4.88-2.56)(0.050+0.150)} = \exp^{-0.464} = 0.63$

reduces the activation cross section of ${}_{8}O^{18}$ to 2.2×10^{-4} barn and the activation cross section for the normal oxygen atom is 4.4×10^{-7} barn. (The abundance of ${}_{8}O^{18}$ in normal oxygen is 0.20 percent.)

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** Now with General Electric Research Laboratories, Schenectady, New York. ¹J. M. Cork, J. R. Richardson, and F. N. D. Kurie, Phys. Rev. 49, 10. March 1998 (2019) (2019 208 (1936).
3F, N. D. Kurie, J. R. Richardson, and H. C. Paxton, Phys. Rev. 49, 368 (1936).
4D be published in the Manhattan Project Technical Series.

 ⁴⁷ Job (1930).
 ⁵ To be published in the Manhattan Project Technical Series.
 ⁴ A survey of all thermal neutron activation cross sections measured at the Argonne Laboratory will be published shortly in *The Physical Project*. Review. ⁵ A barn equals 10⁻²⁴ cm².

Proposal of a Method for the Separation of He³ from He⁴

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THE isotope He³ is present in helium according to Alvarez and Cornog in a concentration 10^{-7} to 10^{-8} . In a discussion with Earl Long about the problem of how to separate this rare He isotope from the bulk of He4, it occurred to the present author that the superfluid state of liquid helium may offer an opportunity to carry out a separation process.

In London's theory the occurrence of superfluidity is explained as a quality acquired at very low temperatures by an atomic species obeying the Bose-Einstein statistic. He⁴ having no spin is supposed to follow this statistic; but for He³, which has odd nuclear spin, the rules of the Fermi statistic are valid and, correspondingly, no superfluid state is expected to occur for this isotope. Therefore, a great enrichment of He³ in liquid He may be achieved by letting the bulk of He4 run out of a container on account of its superfluidity, while He³ is expected to remain in the residue.

If such an experiment should prove to be successful, it would not only have the practical value of producing a supply of He³ to be used for experiments in nuclear physics, but would at the same time offer direct evidence in favor of London's theory.

The drawback of the proposed experiment is the fact that greater quantities of liquid helium (about 10 liters) are needed to carry it out. Since such amounts are usually not available, this proposal is published to bring it to the attention of those laboratories in which the special equipment for the production of greater quantities of liquid He is available.