

the range cell. The range cell was 6.86 cm long with a mica foil on the end away from the target cell. The pressure in the range cell could be varied, allowing a change of about 6 cm in the air equivalent of the range cell. The ionization chamber and foil holder were fastened next to the exit window of the range cell. The entire assembly of target cell, range cell, foil holder and ionization chamber were screwed firmly together so that no relative motion of the parts could take place.

The pressure in the range cell could be read to about 1 mm Hg. The set of mica foils used in this experiment were all from one sheet and the stopping power in mg/cm²/cm of air was measured for two thin foils. The air equivalents of the thicker foils were obtained using the value 1.42₅ mg/cm²/cm air for 5-Mev alpha-particles. The value 1.07 Mev for the energy of the deuterons was computed from the physical constants of the cyclotron. A high bias was used on the counting Thyatron of the amplifier and a correction made for the bias and depth of the ionization chamber using the specific ionization curves for alpha-particles¹ and protons.²

Two proton groups have been measured by Cockcroft and Lewis³ from the N¹⁴(*dp*)N¹⁵ reaction. These proton groups were measured in the present work and, in addition, a group of protons of 66.1-cm range was observed. This 66-cm group was observed both with the N¹⁴-N¹⁵ mixture and with ordinary tank nitrogen in the target cell, but was not observed with CO₂ as the target gas or with the target cell evacuated. Thus there is reason to assign the 66-cm proton group to the N¹⁴(*dp*)N¹⁵ reaction.

With the same N¹⁴-N¹⁵ mixture in the target cell three groups of alphas were observed. Two of these have been found previously^{3, 4} and are from the N¹⁴(*dα*)C¹² reaction. The other group (5.09-cm range) can be attributed to the N¹⁵(*dα*)C¹³ reaction since it appeared with no target gas except the one containing N¹⁵. Further, the *Q* value, 7.40 Mev, checks sufficiently well with the *Q* value 7.55 calculated from the mass values.⁵ In addition to the three homogeneous groups the intense continuous distribution of alphas arising from the reaction N¹⁴+H²→4He⁴ was found below 3.9-cm range.

Because of the relatively large number of alpha-particles in the region below 6 cm it was not possible with the present equipment to detect the short range (~3.5 cm) proton group expected from the N¹⁵(*dp*)N¹⁶ reaction. It is hoped that with a target of much higher N¹⁵ concentration it will be possible to detect protons in this region below 6 cm.

TABLE I. *Experimental results of deuteron bombardment of N¹⁴ and N¹⁵.*

REACTION	RANGE CM 15°, 760 MM	Q VALUE MEV	EXCITATION LEVEL MEV
N ¹⁴ (<i>dα</i>)C ¹²	11.59	13.21	C ¹² gr. st.
	6.37	8.86	4.35
N ¹⁴ (<i>dp</i>)N ¹⁵	90.76	8.41	N ¹⁵ gr. st.
	66.10	6.88	1.53
	20.99	3.10	5.31
N ¹⁵ (<i>dα</i>)C ¹³	5.09	7.40	C ¹³ gr. st.

The results of the present experiment are shown in Table I. The *Q* values obtained from range measurements are all lower by about 0.15 Mev than those obtained from mass values.⁵ Repeated checks on the method used in measuring the ranges indicate that a systematic error is not likely in this part of the experiment. It is possible that there is an error in the energy value 1.07 Mev of the incident deuterons, although this figure was checked by direct measurement of the deuteron ranges.

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A Radioactive Isomer of Sr⁸⁷

In connection with studies soon to be reported of the radioactive isotopes of Sr and Y we have given particular attention to a period of 2.7±0.2 hours which appears to be associated with an excited state of stable Sr⁸⁷.

Stewart, Lawson and Cork¹ reported a 3-hour Sr period produced by Sr+*d* and Sr+*n* (slow). They assigned it to an isomeric state of Sr⁸⁹, rejecting other possible assignments because the activity was electron rather than positron emitting. Cloud-chamber observations showed an apparently continuous spectrum with an upper limit of 610 kev. Our cloud-chamber measurements are similar to theirs, but the beta-ray spectrograph shows that the spectrum consists only of a strong line at *H_p*2340 (360 kev). (The broadening of the spectrum in the cloud chamber is undoubtedly due to the scattering which is important at these low energies.) These electrons are evidently *K* and *L* conversion electrons accompanying a 370-kev gamma-ray emitted in a *K*-electron capture or an isomeric transition. This removes the necessity of assignment to Sr⁸⁹.

We further find that this same period with the same electron spectrum is produced strongly by Rb+*p* at 6 Mev. Only a *p*-*n* reaction could account for the yield, which restricts the assignment to either Sr⁸⁵ or Sr^{87*}. There is also a strong period of ~60 days produced in this reaction. In view of the argument below this may be assigned to Sr⁸⁵.

Finally an 85±5-hour yttrium activity (produced either by Sr+*p* or Sr+*d*) shows the same electron line. When Sr is extracted from an aged sample of this activity the Sr fraction decays with the 2.7-hour period while the Y fraction grows with this period. The only Y isotopes which could be produced by both Sr(*p*, *n*) and Sr(*d*, *n*) are Y⁸⁷ and Y⁸⁸. The latter has already been assigned¹ a 2-hour (*e*⁺) period, since it is produced also by Y+fast neutrons. Stewart *et al.*¹ reported a 60-hour Y period from Sr+*d* which they identified with the 60-70-hour Y⁹⁰, produced by Y+*n* and Y+*d*, postulating a Sr⁸⁸(*d*, *γ*)Y⁹⁰ reaction.

Since the Y decay curves are complicated by the growth of the 2.7-hour Sr and by a very long Y period we believe that they were also observing this 85-hour period. This activity does not show the high energy beta-rays of Y⁹⁰.

We conclude that both the 85-hour and 2.7-hour periods must be assigned to isobars of mass 87. Then Y⁸⁷ decays by K-electron capture to the excited Sr^{87*} which goes to the ground state Sr⁸⁷ with the 2.7-hour period and the emission of a partially converted 370-kev gamma-ray. Absorption measurements in Pb show the expected absorption coefficient for this gamma-ray and Al absorption curves indicate a conversion factor of about 15 percent. The 2.7-hour activity emits x-rays also which show high mass absorption coefficients in both Se and Br indicating they are Sr rather than Rb K x-rays. The same Sr^{87*} is also produced directly by Rb⁸⁷(p, n) and Sr⁸⁶(n, γ). In the Sr+d and Sr+p bombardments it is probably produced only by the decay of Y⁸⁷, which accounts for the low and variable yield.¹

The question arises whether Sr^{87*} is formed by the decay of naturally radioactive Rb⁸⁷. We extracted Sr from 5 g

of RbCl but found no activity as much as 0.1 percent of that expected if every Rb⁸⁷ atom decayed to Sr^{87*}. Furthermore, Libby and Lee² have found the maximum energy of the electrons from Rb⁸⁷ to be 132 kev. There is thus no evidence for the 360-kev electrons from Sr^{87*} and it seems probable that only the decay to the ground state of Sr^{87*} is energetically possible.

It is interesting that in three of the five known pairs of adjacent isobars there is now known an excited radioactive state in one of the pair (In¹¹³ and In¹¹⁵ being the other cases³). In each case the period of the excited state is in the neighborhood of 2-4 hours and the energy is in the range 330-390 kev above the ground state.

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