Neutron Thresholds in Light Elements*†

E. B. TUCKER[‡] Yale University, New Haven, Connecticut (Received May 1, 1951)

A N investigation of the neutron thresholds obtained by deuteron bombardment of lithium, beryllium, boron (natural and enriched¹), carbon, and lead fluoride targets has been carried out. Thresholds have been reported previously at deuteron energies of 920, 990, and 1920 kev²⁻⁴ for the reaction Be⁹(d, n)B¹⁰ and at 1.84 Mev⁴ for O¹⁶(d, n)F¹⁷. A number of (p, n) thresholds have also been investigated.^{4.5}

The apparatus used (Fig. 1) was complicated by the high

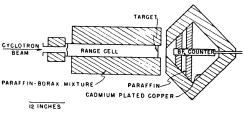


FIG. 1. Apparatus for detection of neutron thresholds.

neutron background and the need for varying the deuteron energy. In addition to the shielding shown in the diagram almost a ton of water was found necessary to cut down the background from the cyclotron, and even so the background level was high. The range cell for varying the bombarding energy was filled with helium. The average values of stopping power of helium needed in calculations were obtained by numerical integration of Mano's results.^{6,7} In order to check on these calculations two experimental evaluations were made of the stopping power of helium relative to air. A direct comparison of deuteron ranges in helium and in air was made, and a value of stopping power was also calculated from the experimental threshold results assuming the previously published value of the lowest $Be^{9}(d, n)B^{10}$ threshold (920 kev). Both of these determinations were slightly higher than the average stopping power calculated from Mano's work but were within the estimated experimental errors.

The experimental curves (Fig. 2) show neutron yield plotted against average deuteron energy, using a value of atomic stopping power of helium relative to air of 0.350. The values assigned to the thresholds are calculated, using the extrapolated beam energy (3.9 Mev) and stopping powers which depend on the energy range concerned (the maximum variation from the value 0.350 is about 2 percent). The probable error is estimated in all cases to be ± 0.15 Mev. The yield curve for carbon shows only resonance peaks, as can be seen by comparison with previous results.⁸ The detection system is such that both resonances and thresholds can be observed. In general, it is expected that the thresholds will show a sharper increase than the resonances, and when using a homogeneous beam the over-all shapes of response should differ considerably. Another method of distinguishing between the two types of yield is to compare fast neutron yield curves with BF³ counter yield curves. The yield obtained from a lithium target (not illustrated) could also be attributed to previously reported resonance levels.⁹ The results obtained for the reaction $Be^{9}(d, n)B^{10}$ indicate two levels. The lower threshold has a value of 0.91 Mev, in good agreement with previous work.^{2-4, 10, 11} The upper inflection point at 2.3 Mev might well be due to a resonance level judging by its sharpness. Ajzenberg¹¹ has obtained levels in B¹⁰ at 6.37, 6.57, and 6.81 Mev but finds them weak compared with levels at 5.91 and 6.11 Mev. Owing to poor resolution (the width of the deuteron beam at half-maximum was 150 kv) a combination of these levels might account for the response observed. Bonner,⁴ however, obtains a threshold at a deuteron energy of 1.920 Mev.

The curve for $F^{19}(d, n)Ne^{20}$ indicates an increase in yield at 1.3 Mev and a steady increase thereafter. No fast neutron yield

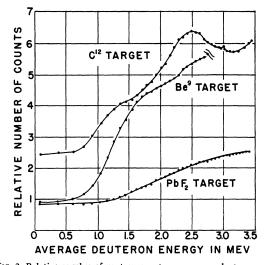


FIG. 2. Relative number of neutron counts versus mean deuteron energy for C¹², Be⁹, and PbF₂ targets. For relative yields from the different targets multiply yield for Be⁹ by 4, PbF₂ by 2. The ordinate then represents hundreds of counts on a scale of 256. The Be⁹ curve has been terminated at 2.7 Mev to prevent confusion. No beryllium thresholds were observed above this energy.

curve has been obtained for this reaction, but the shape of the yield indicates a broad resonance in Ne^{21} about 20 Mev above ground. An exact value cannot be given because no yield maximum is obtained. If it is a true neutron threshold, this would indicate that close spaced levels begin in Ne^{20} at 11.9 Mev above ground (see reference 12 for energy level diagram of Ne^{20}).

Yields from targets of natural and enriched boron (96 percent B¹⁰) are compared in Fig. 3. It appears that the reaction B¹¹(d, n)C¹² is responsible for the entire yield. Results¹³ on the fast neutron yield from B¹¹(d, n)C¹² indicate that the first slow increase (about 1.3 Mev) is due to a resonance level in C¹³. The 1.9-Mev threshold agrees very well with the 1.84 threshold for both targets seems to indicate an oxygen contamination. The rise at

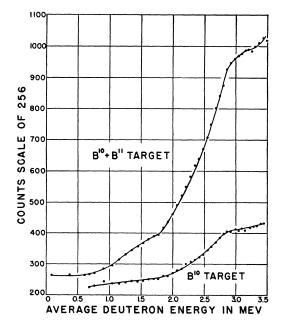


FIG. 3. Neutron yield versus mean deuteron energy for B(d, n)C. The $B^{10} + B^{11}$ target was natural boron (82 percent B^{11}), while the B^{10} target was 96 percent B^{10} .

the end of the curve is not complete enough to obtain an accurate threshold energy, but an approximate value of 3.4 Mev is assigned to it. There is no way of determining whether this last rise is due to a resonance level or not. The fast neutron yield curve13 referred to above does not reach this energy.

The author would like to thank Professor E. C. Pollard for suggesting the problem and for advice during the work. Also the author is grateful to T. W. Bonner for communicating results before publication.

* Supported by the joint program of the ONR and AEC. † Part of a dissertation submitted to the Graduate School of Yale Uni-versity in partial fulfillment of the requirements for the degree of Doctor

I fait of a dissertation submitted to the oradinate School of Yale Oniversity in partial fulfilment of the requirements for the degree of Doctor of Philosophy.
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Alpha-Particle Range-Energy Curve for Kodak NTA Emulsions*

F. E. STEIGERT, E. C. TOOPS, AND M. B. SAMPSON Indiana University, Bloomington, Indiana (Received May 23, 1951)

HE range-energy curves for Ilford B1 emulsions reported by Lattes, Fowler, and Cuer¹ have been used as standards for several years. Recent observations, however, indicate that the alpha-particle curve rises too steeply at the higher energies. Berlman² has modified the proton curve for use with alphaparticles and obtained a curve suitable for Ilford E1 plates. A modified curve has been derived for Kodak NTA emulsions at this laboratory.

The stopping power of the emulsion relative to air for alphaparticles was calculated as a function of the energy by using the procedure outlined by Webb.3 The emulsion is approximated by a homogeneous compound having the composition given by Rotblat.4 The atomic stopping powers used were interpolated from plots of stopping power versus atomic number and energy, as constructed from Bethe's⁵ semi-empirical tabulation. This data was then combined with Bethe's⁶ range curves for air, and the resultant curve of range in emulsion versus alpha-particle energy

TABLE I. Calculated and observed α -particle ranges.

Calculated		Observed		
Alpha- energy Mev	Range microns	Source	Energy Mev	Range microns
2.07	6.88	Po	5.30	21.2
3.00	10.44	ThC	6.06	26.8
4.00	14.50	ThC'	8.78	47.1
5.00	19.6	$Al^{27}(d, \alpha)Mg^{25}$		
6.00	25.6			
7.00	32.2	(0) 124°	13.77 ± 0.08	91.3
8.00	39.5	(I) 90°	13.83	96.6
9.00	47.3	(I) 89°	13.87	97.4
10.00	55.6	(0) 90°	14.34	101.6
11.00	65.0	(I) 60°	14.36	103.9
12.00	74.9	(0) 89°	14.38	103.0
13.00	85.2	(0) 60°	14.87	109.9
14.00	96.4			
15.00	108.7			

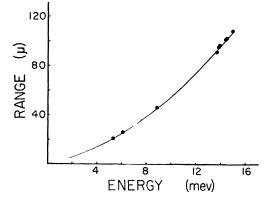


FIG. 1. Calculated alpha-particle range-energy curve for Kodak NTA emulsions, with experimental check points.

shown in Fig. 1 was obtained. Points were calculated at 0.5-Mev intervals, representative values being given in Table I. The curve rise is less steep than indicated by Lattes, Fowler, and Cuer, and there appears to be good agreement with the recent work of Rotblat. The agreement of this curve with experimental data has been obtained in this laboratory, extending the accurate measurement of tracks to 14.9 Mev. Experimental points were taken using natural alpha-particles from polonium and thorium-active deposit sources and the two long-range alpha-groups in the Al²⁷ (d, α) Mg²⁵ reaction, observed at various angles under Bethe's conditions of good geometry. The Q-values⁷ and deuteron energy are known by magnetic analysis to 10 and 40 kev, respectively. The angle of observation was accurate to 15 minutes. Data obtained using this curve agree very well with published work using aluminum foils and counters.8 Curves showing some of the alpha-groups obtained

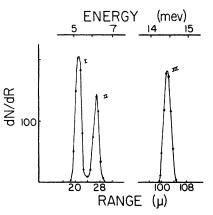


FIG. 2. Typical data obtained from plates. Peak I, Po; II, ThC; III, Al²⁷(d, α)Mg²⁵. The energy scale is plotted above for comparison.

in these experiments are given in Fig. 2. The calibration points on the range-energy curve were obtained by taking the average value from several plates to obtain good statistics. The position of the peaks was observed not to shift as a function of time of exposure in vacuum over the range of several minutes to one hour.

* This work was assisted by the joint program of the ONR and AEC.
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