Eq. (53) assumes the form

$$\langle L_{\alpha}' L_{\beta}' \rangle \sim \operatorname{cosec}^{2} \theta \int_{x, y} \int y(x + y \operatorname{cot} \theta) \\ \times \nabla^{2} \nabla^{2} N(x, y, z) dx dy.$$
 (61)

It can be shown that the expression which remains under the integral sign is changed but little if the circular sector is expanded into a semicircle. Thus, the expression (57) can be further simplified into

$$\langle L_{\alpha}' L_{\beta}' \rangle \sim \frac{1}{2\theta^2} \int_{x, y \to -\infty}^{\infty} xy \nabla^2 \nabla^2 N dx dy + \frac{1}{2\theta^3} \int_{x, y \to -\infty}^{\infty} y^2 \nabla^2 \nabla^2 N dx dy. \quad (62)$$

Of these two terms, the second one is large compared with the first, so that the final expression is

$$\langle L_{\alpha}' L_{\beta}' \rangle \sim \frac{1}{2\theta^3} \int_{x, y=-\infty}^{\infty} y^2 \nabla^2 \nabla^2 N dx dy,$$
 (63)

where y is the direction perpendicular to \mathbf{R} in the plane of source and receivers.

To obtain the usual correlation coefficient, the expression (63) has to be divided through by (50). It is found that the correlation coefficient is independent of the distance between source and receivers and inversely proportional to the cube of the distance between the two receiving stations perpendicular to the line connecting the source with the location of the receivers, δ , as indicated in Fig. 2.

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A Calculation of the Binding Energies of H^3 and He^4 with a New Potential

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The binding energies of the nuclei H³ and He⁴ are calculated by the method of equivalent two-body, using the potential suggested by Wang. The range at which the potential between two nucleons is cut off is the same as that for the case of the deuteron, and the same range for the equivalent two-body is deduced accordingly. The binding energies thus calculated are 7.3 Mev and 15.1 Mev, respectively.

I. METHOD OF CALCULATION

`HE nuclear potential proposed by Wang¹ was previously applied by the author to the calculation of the binding energy of the deuteron and of the neutron-proton scattering.² In the present work the binding energies of the nuclei H³ and He⁴ are computed by using the same potential in the method of the equivalent two-body.

Let $C\phi(r)$ be the potential between any two nucleons. If the Gaussian wave function $N \exp[-\frac{1}{2}\nu(r_{12}^2 + r_{13}^2 + r_{23}^2)]$ is used, the variation energy for H³ is³

$$E(H^{3}) = \frac{9\nu\hbar^{2}}{2M} - 3C\frac{4}{\pi^{\frac{1}{2}}} \int_{0}^{\infty} \exp(-x^{2}) \times \phi[x/(3\nu/2)^{\frac{1}{2}}]x^{2}dx, \quad (1)$$

where

where

$$x = (3\nu/2)^{\frac{1}{2}}r_{12}$$
 (or r_{13}, r_{23}),

and r_{12} is the distance between particles 1 and 2. For the nucleus He⁴, taking $N \exp\left[-\frac{1}{2}\nu(r_{12}^2+r_{13}^2+r$ $+r_{14}^2+r_{23}^2+r_{24}^2+r_{34}^2)$] as the wave function, we have

$$E(\text{He}^{4}) = \frac{9\nu\hbar^{2}}{M} - 6C\frac{4}{\pi^{\frac{1}{2}}} \int_{0}^{\infty} \exp(-x^{2}) \times \phi[x/(2\nu)^{\frac{1}{2}}]x^{2}dx, \quad (2)$$

³ William Rarita and R. D. Present, Phys. Rev. 51, 788 (1937).

¹ K. C. Wang and H. L. Tsao, Phys. Rev. 66, 155 (1944); Nature 155, April 28 (1945).

² Mu-Hsien Wang, Phys. Rev. 66, 103 (1944).

5.00×10 ⁻						
ε, erg	<u>2</u> ε	$\coth (370\epsilon^{\frac{1}{2}})$				
10.8×10 ⁻⁶	1.32	1.20-				
11.5×10-6	1.17	1.18				
11.7×10-6	1.14	1.17				

TABLE II.

5.00×10 ⁻⁵						
e, erg	2e	$\cosh(388\epsilon^{\frac{1}{2}})$				
11.5×10-6	1.18	1.16				
11.6×10-6	1,16	1.15+				
12.0×10 ⁻⁶	1.08	1.15				

where

$$x = (2\nu)^{\frac{1}{2}} r_{12}$$
 (or r_{13}, r_{14}, \cdots).

Now, let $D\Phi(r)$ be the potential for the equivalent two-body problem. Using the wave function $N \exp[-\mu r^2/2]$, we get for the variation energy for the equivalent two-body:

$$E_{\rm eq} = \frac{3\mu\hbar^2}{2M} - D\frac{4}{\pi^{\frac{1}{2}}} \int_0^\infty \exp\left(-x^2\right) \\ \times \Phi(x/\mu^{\frac{1}{2}}) x^2 dx, \quad (3)$$
where

 $x=\mu^{\frac{1}{2}}r.$

In order that $E(H^3)$, as given by (1), may equal E_{eq} , we should have

$$\mu = 3\nu, \qquad D = 3C, \qquad \Phi(r) = \phi(2^{\frac{1}{2}}r). \qquad (4)$$

Similarly, to have $E(\text{He}^4) = E_{eq}$, we must set

$$3\mu/2 = 9\nu, \quad D = 6C, \quad \Phi(r) = \phi(3^{\frac{1}{2}}r).$$
 (5)

In our case, the potential is

$$C\phi(r) = V_0 = \text{const. for } r \le a,$$

$$C\phi(r) = -Ae^{K/r} \quad \text{or} \quad -\frac{B}{r}e^{K/r} \quad \text{for } r > a,$$

where

$$r = r_{12}$$
 (or r_{13}, r_{14}, \cdots),

and where a has the same meaning as in the deuteron; its value was found in reference 2. To determine the cut-off distance, r_0 , to be used in the equivalent two-body method, we compare Eq. (3) in turn with Eqs. (1) and (2), and find that

$$r_0 = a/2^{\frac{1}{2}}$$
 for H³, (6)

$$r_0 = a/3^{\frac{1}{2}}$$
 for He⁴. (7)

If the zero cut-off method is employed, the potential for H³ has therefore the form (9a, b) or the form (10a, b), and the potential for He⁴ has the form (11a, b) or the form (12a, b).

The binding energy, ϵ , of H³ and of He⁴ is then the proper value of the wave equation

$$M\Delta\psi - \hbar^2(\epsilon + V)\psi = 0,$$

where V is the potential. The condition for the smooth joint of the wave function at r_0 is²

$$|V(r_0)|/2\epsilon - 1 = \coth(\beta r_0), \qquad (8)$$

where $\beta = (M\epsilon/\hbar^2)^{\frac{1}{2}}$.

II. THE BINDING ENERGY OF H³

 $D\Phi(r) = 0$ for $r \le r_0 = a/2^{\frac{1}{2}}$,

(a) For the potential

and

and

$$D\Phi(\mathbf{r}) = -3A \exp(K/2^{\frac{1}{2}}\mathbf{r}) \text{ for } \mathbf{r} > \mathbf{r}_0, \quad (9b)$$

where² $A = 4.78 \times 10^{-45}$, $K = 3.84 \times 10^{-11}$ cm, $a = 4.21 \times 10^{-13}$ cm, Eq. (8) becomes:

 $5.00 \times 10^{-5}/2\epsilon - 1 = \operatorname{coth}(370\epsilon^{\frac{1}{2}}).$

The numerical results are given in Table I. The estimated binding energy of H³ is therefore 11.5×10^{-6} erg, or 7.23 Mev.

(b) For the potential

$$D\Phi(r) = 0$$
 for $r \le r_0 = a/2^{\frac{1}{2}}$, (10a)

$$D\Phi(r) = -\frac{3B}{2^{\frac{1}{2}}r} \exp(K/2^{\frac{1}{2}}r) \quad \text{for } r > r_0, \quad (10b)$$

where² $a = 4.42 \times 10^{-13}$ cm, $B = 1.84 \times 10^{-55}$, we have

$$5.00 \times 10^{-5}/2\epsilon - 1 = \coth(388\epsilon^{\frac{1}{3}})$$

The results are given in Table II. The estimated binding energy of H³ is therefore 11.6×10^{-6} erg, or 7.30 Mev. These results are in good agreement with the experimental value 8.3 Mev.

III. THE BINDING ENERGY OF He⁴

(a) For the potential

$$D\Phi(r) = 0 \text{ for } r \le r_0 = a/3^{\frac{1}{2}},$$
 (11a)

(9a)

TABLE III		TABLE IV		
1.00×10-4		1.00×10-4		
<u></u> 2e	$ \coth (302\epsilon^{\frac{1}{2}}) $	e, erg	<u></u> -1 2e	$\coth (317\epsilon^{\frac{1}{2}})$
1.12	1.12-	23.8×10^{-6}	1.11	1.10-
1.11	1.11	24.0×10^{-6} 24.2 × 10^{-6}	1.08	1.09
		TABLE III $\frac{1.00 \times 10^{-4}}{2\epsilon} - 1$ $\coth (302\epsilon^{\frac{1}{2}})$ 1.12 1.12^{-1} 1.11 1.11 1.08 1.10	TABLE III $\frac{1.00 \times 10^{-4}}{2\epsilon} - 1$ coth $(302\epsilon^{\frac{1}{9}})$ 1.12 1.12^{-} 23.8×10^{-6} 1.11 1.11 24.0×10^{-6} 1.08 1.10 24.2×10^{-6}	TABLE III TABLE IV $\frac{1.00 \times 10^{-4}}{2\epsilon} - 1$ $coth (302\epsilon^{\frac{1}{2}})$ ϵ, erg $\frac{1.00 \times 10^{-4}}{2\epsilon} - 1$ 1.12 $1.12^ 23.8 \times 10^{-6}$ 1.11 1.08 1.10 24.2×10^{-6} 1.07

and

$$D\Phi(r) = -6A \exp(K/3^{\frac{1}{2}}r)$$
 for $r > r_0$, (11b)

where the constants have the same values as in Section II (a), Eq. (8) turns out to be

$$1.00 \times 10^{-4}/2\epsilon - 1 = \coth(302\epsilon^{\frac{1}{2}}),$$

and yields Table III. According to this table, the binding energy of He⁴ is 23.8×10^{-6} erg, or 15.0 Mev.

(b) For the potential

$$D\Phi(r) = 0$$
 for $r \le r_0 = a/3^{\frac{1}{2}}$, (12a)

and

$$D\Phi(\mathbf{r}) = -\frac{6B}{3^{\frac{3}{2}}r} \exp(K/3^{\frac{3}{2}}r) \quad \text{for } \mathbf{r} > r_0, \quad (12b)$$

where the constants have the same values as in Section II (b), Eq. (8) becomes

$$1.00 \times 10^{-4}/2\epsilon - 1 = \coth(317\epsilon^{\frac{1}{2}})$$

and yields Table IV. According to this table, the binding energy of He⁴ is 24.0×10^{-6} erg, or 15.1 Mev.

The values of the binding energy of He⁴ computed above are about 45 percent less than the experimental value 27.8 Mev. The results for He^4 are therefore not as good as those for H^3 . This may mean that the method of the equivalent two-body tends to be a poor approximation as the number of particles increases.

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Recurrence Phenomena in Small Cosmic-Ray Bursts*

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Chree's method of superposed epochs was employed in the statistical investigation of variations in the frequency of occurrence of small cosmic-ray bursts which produced 2.9 to 3.6 millions of pairs of ions in a shielded spherical ionization chamber of 13.3 liters effective volume containing air at 160 atmospheres. The data employed were obtained by Long and Whaley in the same investigation (during a little more than 18 months in 1938 and 1939) which supplied the data for the author's work on recurrences in variations of cosmic-ray intensity and their relation to geomagnetic and heliophysical activities. The

HE data regarding bursts which are discussed in this paper were observed by Long¹ and Whaley^{2,3} in the same investigation

analysis was carried out only for the range of day numbers from -45 to +45. Irregular secondary pulses were found both preceding and subsequent to both positive and negative primary pulses. Both subsequent and preceding difference curves and the combination difference curve displayed secondary pulses with peaks at about 27 days preceding and subsequent to the primary pulses. The secondary pulses amounted to about 3 or 4 percent of the average frequency of approximately 37 small bursts per day, and about 10 percent of the larger variations constituting the primary pulses.

during eighteen months in 1938 and 1939, which vielded data used heretofore by the writer.4,5

^{*} Presented at the meeting of the American Physical Society at St. Louis, November 30-December 1, 1945; Phys. Rev. 69, 46 (1946). ¹V. A. Long, Ph.D. Thesis, University of Colorado, August 14, 1940.

² R. M. Whaley, M.A. Thesis, University of Colorado, June 3, 1940.

⁸V. A. Long and R. M. Whaley, Phys. Rev. 59, 470 (1941). ⁴ J. W. Broxon, Phys. Rev. **59**, 773 (1941). ⁵ J. W. Broxon, Phys. Rev. **62**, 508 (1942).