TERMINATION OF THE PROTON-PROTON CHAIN IN STELLAR INTERIORS*

Peter D. Parker,[†] John N. Bahcall, and William A. Fowler

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ABSTRACT

The roles of the various H²-burning and He³-burning reactions in the termination of the protonproton chain in stellar interiors have been re-examined using (1) recent experimental determinations of the cross-sections for the H²(p, γ)He³ and He³(α , γ)Be⁷ reactions, (2) theoretical studies of the rates of the relevant beta-decay reactions, and (3) a re-examination of the previous cross-section factor determinations for the other possible reactions in the proton-proton chain.

I. INTRODUCTION

A number of review articles have discussed the termination of the proton-proton chain, for example, Bethe (1939), Salpeter (1952), Burbidge, Burbidge, Fowler, and Hoyle (1957), and Fowler (1960). However, recent experimental studies of the $H^2(p, \gamma)He^3$ reaction by Griffiths, Lal, and Scarfe (1963) and recent theoretical and experimental studies of the termination of the chain via the $He^3(a, \gamma)Be^7$ reaction (Kavanagh 1960; Bahcall 1962b; Parker and Kavanagh 1963) make it desirable to reexamine the roles of various reactions in the termination of the proton-proton chain. In the course of this re-examination it has been found that, contrary to statements in some previous review articles, further experimental investigation of the Li⁴ system is necessary before one can definitely rule out the possibility that Li⁴ plays an important role in the termination of the proton-proton chain.

In the calculations described below we have adopted many of the definitions and notations of Burbidge *et al.* (1957). The cross-section factor, S(E), is defined by the following equation:

$$S(E) = \sigma(E)E \exp \left(\frac{31.28 Z_1 Z_0 A^{1/2}}{E^{1/2}} \right) \text{ keV-barns}, \tag{1}$$

where σ is the reaction cross-section in barns, E is the center-of-mass energy in keV, Z_1 and Z_0 are the charges of the interacting nuclei in units of the proton charge, and A is the reduced mass, $A_1A_0/(A_1 + A_0)$, in atomic mass units. The mean reaction rate, P, of a non-resonant nuclear reaction can be written

$$P = 7.20 \times 10^{-19} n(1) n(0) f S \tau^2 e^{-\tau} (1 + \delta_{10})^{-1} (A Z_1 Z_0)^{-1} \text{ reactions } \text{cm}^{-3} \text{ sec}^{-1}, \qquad (2a)$$

$$P = 2.62 \times 10^{29} \rho^2 \frac{X_1 X_0}{A_1 A_0} fS \tau^2 e^{-\tau} (1 + \delta_{10})^{-1} (AZ_1 Z_0)^{-1} \text{ reactions cm}^{-3} \text{ sec}^{-1}, \quad \text{(2b)}$$

where

$$\tau = 42.48 \left(Z_1^2 Z_0^2 \frac{A}{T_6} \right)^{1/3}.$$
 (2c)

In equations (2a), (2b), and (2c) T_6 is the temperature in millions of degrees; δ_{ij} is the Kronecker delta,

$$\delta_{ij} = 1$$
 if $i = j$, $\delta_{ij} = 0$ if $i \neq j$;

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[†] Present address: Brookhaven National Laboratory, Upton, New York.

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 ρ is the density in gm cm⁻³; n(i) is the number density in particles per cm³ of nuclei of type (i); X_i is the fraction by weight of nuclei of type (i) in the medium,

$$n(i) = 6.023 \times 10^{23} \rho \, \frac{X_i}{A_i}; \tag{3}$$

S is the cross-section factor in keV-barns defined in equation (1); and f is the electronscreening factor discussed by Salpeter (1954). In the weak-screening approximation appropriate for the sun, Salpeter has shown that

$$f = \exp\left(0.188 \, Z_1 Z_0 \zeta \rho^{1/2} T_6^{-3/2}\right),\tag{4a}$$

where

$$\zeta = \left[\sum_{i} \left(X_{i} \frac{Z_{i}^{2}}{A_{i}} + X_{i} \frac{Z_{i}}{A_{i}}\right)\right]^{1/2}.$$
(4b)

It is convenient to introduce a quantity, $\langle i, j \rangle$, defined by the relations

$$\langle i, j \rangle \equiv \frac{P(1 + \delta_{i, j})}{n(i)n(j)} \, \mathrm{cm}^{3} \, \mathrm{sec}^{-1},$$

$$\langle i, j \rangle = \langle \sigma v \rangle_{ij}.$$

$$\langle i, j \rangle = \langle j, i \rangle,$$

$$(5)$$

where $\langle i, j \rangle$ is the thermal average of the cross-section times the relative velocity of particles *i* and *j*. Using equation (5) we can write the mean lifetime of a nucleus (*i*) for interaction with nuclei of type (*j*) as follows:

$$\tau_j(i) = [\langle i, j \rangle n(j)]^{-1} \sec .$$
⁽⁶⁾

The rate, $R_j(i)$, at which nuclei of type (i) are consumed by interaction with nuclei of type (j) can be written

$$R_{j}(i) = P(1 + \delta_{i, j})$$

= $\langle i, j \rangle n(i) n(j)$ nuclei cm⁻³ sec⁻¹. (7)

The fraction of the nuclei (i) destroyed that are consumed by interaction with nuclei (j) is thus

$$F_j(i) = \frac{R_j(i)}{\sum_k R_k(i)},$$
(8)

and the fraction of the nuclei (i) produced that are consumed by interaction with nuclei (j) is

$$\Phi_j(i) = \frac{R_j(i)}{\text{Rate of production of }(i)}.$$
(9)

Under equilibrium conditions the rates of production and consumption of nuclei (i) are equal, and therefore

$$F_j(i) = \Phi_j(i) . \tag{10}$$

The fraction of the He⁴ that is produced by the interaction of nuclei (i) and (j) is defined as

$$C_{\text{He}^{4}}(i, j) = \frac{1}{(1+\delta_{ij})} \frac{\langle i, j \rangle n(i) n(j)}{\text{Rate of production of He}^{4}}$$

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The cross-sections for the nuclear reactions that occur in the proton-proton chain cannot in general be measured at the energies of interest to astrophysicists (of the order of 20 keV). Hence, for the cross-section factor, extrapolations from higher-energy measurements must be used to obtain a zero-energy intercept, S_0 , and an average value for the derivative at low energies, $(dS/dE)_0$. Note that $S_0 = S(E = 0)$ is not identical with the $S_o = S(E = E_o)$ of Burbidge, Burbidge, Fowler, and Hoyle (1957). Christy and Duck (1961) and Tombrello and Parker (1963) have discussed the use of direct-capture calculations to obtain reliable extrapolations from high energy data. The Q-value and the values of S_0 and $(dS/dE)_0$ for the various reactions in the proton-proton chain are listed in Table 1. Estimates of the absolute errors involved in the determination of S_0 are included for those cases where it was felt that such estimates were meaningful. In the calculations described below, we have followed Caughlan and Fowler (1962) and have included the temperature dependence of S by writing $S(T_6)$ as

$$S(T_6) = S_0 \left\{ 1 + \frac{5}{12\beta} T_6^{1/3} + \frac{(dS/dE)_0}{S_0} [E_0(T_6) + \frac{35}{36}kT_6] + \dots \right\}, \qquad (11a)$$

where

$$\beta = 42.48 \ (Z_1^2 Z_0^2 A)^{1/3} \tag{11b}$$

and

$$E_0(T_6) = 1.220 \ (Z_1^2 Z_0^2 A T_6^2)^{1/3} \text{ keV} . \tag{11c}$$

II. NUCLEAR ABUNDANCES

In the proton-proton chain, unlike the CNO bi-cycle, many nuclear reactions occur in which neither of the interacting nuclei is a proton. Furthermore, the interacting nuclei have abundances which are not independent of each other but are instead determined by the physical conditions in the star in which the reactions are occurring. Because many reactions interrelate the various nuclear species, there is no simple way of expressing the abundance of each of these species in terms of the proton abundance, $X_{\rm H}$. Hence, it is not possible to express our results for the pertinent reactions in the form of a simple table of [log ($\rho \tau X_{\rm H}/100$)] versus T_6 , as Caughlan and Fowler (1962) have done for the CNO bi-cycle. Instead, in the discussion that follows, we present general expressions which can be used to calculate the quantities of interest and actually carry out such calculations (except where noted) only for the case of $\rho = 100$ gm cm⁻³ and $X_{\rm H} = X_{\rm He} = 0.50$.

The proton and alpha-particle abundances are described by the parameters $X_{\rm H}$ and $X_{\rm He}$ defined above. The electron density is given by the usual expression,

$$n(e) \simeq 6.023 \times 10^{23} \rho \frac{(1+X_{\rm H})}{2} \,{\rm cm}^{-3}.$$
 (12)

In order to calculate the abundances of deuterium and He³, it is necessary to consider the various reactions involved in the proton-proton chain. The discussions in Sections III and IV will show that the following reactions are the important ones for producing and destroying deuterium and He³ under normal stellar conditions:

(a)
$$H^{1}(p, \beta^{+} \nu)H^{2}$$
, (d) $He^{3}(d, p)He^{4}$,
(b) $H^{2}(p, \gamma)He^{3}$, (e) $He^{3}(\tau, 2p)He^{4}$, (13)
(c) $H^{2}(\tau, p)He^{4}$, (f) $He^{3}(a, \gamma)Be^{7}$,

where we have followed the convention that $\tau \equiv \text{He}^3$ inside the parentheses in the same way that $p \equiv \text{H}^1$, $d \equiv \text{H}^2$, $t \equiv \text{H}^3$, and $a \equiv \text{He}^4$. Thus, it should be noted that reactions (13c) and (13d) are identical except that (13c) is written so as to emphasize its role as

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TABLE 1

REACTION PARAMETERS (PROTON-PROTON CHAIN)

Reaction		Q-Value*	S	$(dS/dE)_0$	$r_j(i)$	Reference
		(JMLEV)	(Kev-Darns)	(Darns)	(yr.)	
$\mathrm{H}^{1}(b, \mathcal{B}^{+}_{p})\mathrm{H}^{2}$		+ 1.442	$(3.36\pm0.4)\times10^{-22}$	$+ 2.7 \times 10^{-24}$	8.85×10^{9}	Fowler (1960)
$H^2(p, \gamma)He^3$	•	+ 5 493	$(2.5 \pm 0.4) \times 10^{-4}$	$+ 7.9 \times 10^{-6}$	4.43×10^{-8}	Griffiths et al. (1963)
$H^2(d, p)H^3$		+ 4.032	100	+ 0.525	5.32×10^{5}	Arnold <i>et al.</i> (1954)
$H^{2}(\sigma, n)He^{\circ}$		± 3.200 ± 18350	67×10 ³	+27	1.15×10^{-4}	Arnold <i>et al.</i> (1954)
$\operatorname{He}^{3}(d, b)\operatorname{He}^{4}$		+18.352	6.7×10^{3}	+27	8.86×10^{8}	Arnold et al. (1954)
$He^{3}(\tau, 2b)He^{4}$		+12.859	1.1×10^{3}	-	5.28×10^{5}	Good et al. (1954)
$He^{3}(a, \gamma)Be^{7}$	•	+1.586	(0.47 ± 0.05)	-2.8×10^{-4}	970×10^{6}	Parker et al. (1963)
$\operatorname{Be}^{7(e^{-}, v)}\operatorname{Li}^{7}$:	+ 0.861 (89 7%)		-	2 86 V 10-1	\int Bahcall (1962b)
$\operatorname{Be}^{7}(e^{-}, v)\operatorname{Li}^{7*}$	•	+ 0.383(10.3%)				Taylor et al. (1962)
$\mathrm{Li}^{7}(p, \alpha)\mathrm{He}^{4}$		+17.347	120	0.0	1.80×10^{-5}	Sawyer and Philips (1953)
$\text{Li}^7(b, \gamma) \text{Be}^8(a) \text{He}^4$	•	+17.347	~1	•	$>2 \times 10^{-3}$	Sawyer and Philips (1953)
$\operatorname{Be}^{r(p, \gamma)} B^{8}$		+ 0.135	(0.030 ± 0.01)	0	8.81×10^{10}	Kavanagh (1960)
$B^{8}(\beta^{+}\nu)Be^{8*}(\alpha)He^{4}$		+18.074	•	•	3×10^{-8}	Bahcall $(1962a)$
$\operatorname{He}^{3}(e^{-},\nu)\operatorname{H}^{3}$	•	- 0.018	•		2×10^{10}	Bahcall and Wolf (1963)
$H^{3}(\dot{\rho}, \dot{\gamma})\dot{H}e^{4}$	•	+19.813	1.5×10^{-3}	$+ 2.7 \times 10^{-5}$	1.68×10^{-8}	Perry and Bame (1955)
$\operatorname{He}^{3}(p, \beta^{+}\nu)\operatorname{He}^{4}$	•	+19795	•	•		
* König, Mattauch a	und Waps	tra (1962).	+ τ calcı	ulated for $X_{\rm H} = X_{\rm He} =$	$0.5, \rho = 100 \text{ gm cm}$	m^{-3} and $T_6 = 15$.

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a deuterium-burning reaction and (13d) is written so as to emphasize its role as a He³ burning reaction.

Assuming that reactions (13a)-(13f) are the only important ones, we can write

$$\dot{n}(\mathrm{H}^2) = \frac{1}{2} \langle \mathrm{H}^1, \mathrm{H}^1
angle n(\mathrm{H}^1) n(\mathrm{H}^1) - \langle \mathrm{H}^2, \mathrm{H}^1
angle n(\mathrm{H}^2) n(\mathrm{H}^1) - \langle \mathrm{H}^2, \mathrm{He}^3
angle n(\mathrm{H}^2) n(\mathrm{He}^3)$$
 (14a)

and

$$\dot{n}(\mathrm{He}^{3}) = \langle \mathrm{H}^{2}, \mathrm{H}^{1} \rangle n(\mathrm{H}^{2}) n(\mathrm{H}^{1}) - \langle \mathrm{He}^{3}, \mathrm{H}^{2} \rangle n(\mathrm{He}^{3}) n(\mathrm{He}^{3}) - \langle \mathrm{He}^{3}, \mathrm{He}^{3} \rangle n(\mathrm{He}^{3}) n(\mathrm{He}^{3}) - \langle \mathrm{He}^{3}, \mathrm{He}^{4} \rangle n(\mathrm{He}^{3}) n(\mathrm{He}^{4}) .$$
(14b)

Over a wide range of stellar temperatures the rates of production and consumption of H^2 and He^3 are equal (see Secs. III*a* and IV*a*, respectively) so that

$$\dot{n}(\mathrm{H}^2) = 0, \qquad (15a)$$

and

$$\dot{n}(\mathrm{He}^3) = 0. \tag{15b}$$

Let II = $n(H^2)/n(H^1)$, III = $n(He^3)/n(H^1)$, and IV = $n(He^4)/n(H^1)$. Then under the equilibrium conditions defined by equations (15*a*) and (15*b*) equations (14*a*) and (14*b*) can be rewritten as

$$\frac{1}{2}\langle H^1, H^1 \rangle - \langle H^2, H^1 \rangle II - \langle H^2, He^3 \rangle II III = 0, \qquad (16a)$$

and

$$\langle H^2, H^1 \rangle II - \langle He^3, H^2 \rangle III II - \langle He^3, He^3 \rangle III III - \langle He^3, He^4 \rangle III IV = 0$$
. (16b)

Combining equations (16a) and (16b), we obtain

$$II = \frac{1}{2} \langle H^{1}, H^{1} \rangle / (\langle H^{2}, H^{1} \rangle + \langle H^{2}, He^{3} \rangle III), \qquad (17a)$$

and

$$\begin{array}{l} \langle \mathrm{He^3, H^2} \rangle \langle \mathrm{He^3, He^3} \rangle \mathrm{III^3} + [\langle \mathrm{He^3, He^3} \rangle \langle \mathrm{H^2, H^1} \rangle + \langle \mathrm{He^3, H^2} \rangle \langle \mathrm{He^3, He^4} \rangle \mathrm{IV}] \, \mathrm{III^2} \\ + [\frac{1}{2} \langle \mathrm{He^3, H^2} \rangle \langle \mathrm{H^1, H^1} \rangle + \langle \mathrm{H^2, H^1} \rangle \langle \mathrm{He^3, He^4} \rangle \mathrm{IV}] \, \mathrm{III} - \frac{1}{2} \langle \mathrm{H^2, H^1} \rangle \langle \mathrm{H^1, H^1} \rangle = 0 \,. \end{array} \right.$$

The coupled equations (17*a*) and (17*b*) were solved as a function of temperature for the case $X_{\rm H} = X_{\rm He} = 0.50$, that is, IV = 0.25, and the resulting $n({\rm H}^2)/n({\rm H}^1)$ and $n({\rm He}^3)/n({\rm H}^1)$ ratios are plotted in Figures 1 and 2. It should be noted that except for screening corrections and except under the special conditions described in Section IVf $n({\rm H}^2)/n({\rm H}^1)$ and $n({\rm He}^3)/n({\rm H}^1)$ are independent of density. Using the equilibrium abundances shown in Figures 1 and 2 for H² and He³, for the case $X_{\rm H} = X_{\rm He} = 0.50$ and $\rho = 100$ gm cm⁻³, we have calculated the mean lifetimes (defined by eq. [5]) associ ated with each of the reactions in the proton-proton chain; the results are plotted in Figures 3, A, and B.

III. DEUTERIUM-BURNING REACTIONS

The $H^2(p, \gamma)He^3$ reaction is the only reaction usually discussed in connection with deuterium burning in the interiors of stars operating on the proton-chain. The discussion in this section shows that, in fact, the $H^2(p, \gamma)He^3$ reaction is the most important deu terium-burning reaction in stellar interiors under normal conditions. There are, how ever, other deuterium-burning reactions, and we discuss them in order to determin under what, if any, conditions they contribute significantly to the consumption o deuterium in the proton-proton chain.

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a) $H^2(p, \gamma)He^3$

Griffiths *et al.* (1963) have measured the cross-section for the $H^2(p, \gamma)He^3$ reaction in the energy region from 16 to 35 keV and have obtained the following intercept parameters for S(E):

and

 $S_0 = (0.25 \pm 0.04) \times 10^{-3} \text{ keV-barns}$,

 $(dS/dE)_0 = +0.79 \times 10^{-5}$ barns.



FIG. 1.—The ratio, $n(H^2)/n(H^1)$, of the deuterium equilibrium abundance to the proton abundance is shown for $X_H = X_{H_0} = 0.5$ as a function of T_6 , the temperature in millions of degrees.



FIG. 2.—The ratio, $n(\text{He}^3)/n(\text{H}^1)$, of the He³ equilibrium abundance to the proton abundance is shown for $X_{\rm H} = X_{\rm Ho} = 0.5$ as a function of T_6 . The dashed curve below $T_6 = 6$ indicates that in this temperature region the actual He³ abundance will probably be less than the equilibrium value plotted here.

A comparison of the mean lives for the $H^2(p, \gamma)He^3$ reaction and the $H^1(p, \beta^+ \nu)H^2$ reaction (see Fig. 3, A) shows that the $H^2(p, \gamma)He^3$ reaction is sufficiently rapid at temperatures above a few hundred thousand degrees to guarantee that the rate of consumption of H^2 via the $H^2(p, \gamma)He^3$ reaction will be equal to the rate of production of H^2 via the $H^1(p, \beta^+ \nu)H^2$ reaction. Hence, the assumption that $\dot{n}(H^2) = 0$ is justified at such temperatures, and the abundance of deuterium equals its equilibrium value.

b) $H^2(d, p)H^3$ and $H^2(d, n)He^3$

The cross-sections for these reactions have been measured by Arnold, Phillips, Sawyer, Stovall, and Tuck (1954) over a deuteron-energy range from 13 to 113 keV. Summing their cross-sections for these two reactions, one gets the following intercept parameters for the total S(E):

$$S_0 = 100 \text{ keV-barns}$$
,

and

$$(dS/dE)_0 = +0.525$$
 barns

A comparison of the mean lives for the $H^2 + H^2$ reactions and the $H^1(p, \beta^+ \nu)H^2$ reaction (see Fig. 3, A) shows that the $H^2 + H^2$ reactions are also sufficiently fast to guarantee an equilibrium abundance for deuterium. Moreover, a comparison of the cross-section factors for the $H^2 + H^2$ reactions and the $H^2(p, \gamma)He^3$ reaction shows that the burning of deuterium by deuterium has a considerable advantage over the $H^2(p, \gamma)He^3$ reaction because of the much larger cross-section factor for the $H^2 + H^2$ reaction. However, the relative deuterium-to-proton abundance, which (Fig. 1) is $\leq 10^{-17}$ for $T_6 \geq 1$, completely



FIG. 3.—A, The mean lives of deuterium for the various H²-burning reactions are compared to the mean life of the H¹(ρ , $\beta^+\nu$)H² reaction; all mean lives are plotted as a function of T_6 for $X_{\rm H} = X_{\rm He} = 0.5$ and $\rho = 100$ gm cm⁻³. The dashed curve has the same significance as in Fig. 2. B, The mean lives of He³ for the various He³-burning reactions and the mean lives associated with the various A = 7 products of the He³(α , γ)Be⁷ termination are compared to the mean life of the H¹(ρ , $\beta^+\nu$)H² reaction. All mean lives are plotted as functions of T_6 for $X_{\rm H} = X_{\rm He} = 0.5$ and $\rho = 100$ gm cm⁻³. The dashed curve has the same significance as in Fig. 2.

overcomes this advantage. A calculation of the fraction of deuterium burned by both the $H^2 + H^2$ reactions,

$$F_{d}(\mathrm{H}^{2}) = \frac{\langle \mathrm{H}^{2}, \mathrm{H}^{2} \rangle \mathrm{II}}{\langle \mathrm{H}^{2}, \mathrm{H}^{1} \rangle + \langle \mathrm{H}^{2}, \mathrm{H}^{2} \rangle \mathrm{II} + \langle \mathrm{H}^{2}, \mathrm{He}^{3} \rangle \mathrm{III}},$$
(18)

assuming an equilibrium abundance of He³ and $X_{\rm H} = X_{\rm He} = 0.5$, shows that this fraction is less than 8.41×10^{-14} at all temperatures. Thus, the H² + H² reactions can be ignored in stellar-interior calculations.

c) $H^{2}(\tau, p)He^{4}$

Arnold *et al.* (1954) have measured the cross-sections for the $H^2(\tau, p)He^4$ reaction over a deuteron-energy range from 36 to 93 keV. From their measurements one can obtain the following intercept parameters:

$$S_0 = 6700 \text{ keV-barns}$$
,

and

$$(dS/dE)_0 = +27$$
 barns.

The fraction of deuterium burned by this reaction,

$$F_{\tau}(\mathrm{H}^{2}) = \frac{\langle \mathrm{H}^{2}, \mathrm{H}e^{3} \rangle \mathrm{III}}{\langle \mathrm{H}^{2}, \mathrm{H}^{1} \rangle + \langle \mathrm{H}^{2}, \mathrm{H}^{2} \rangle \mathrm{II} + \langle \mathrm{H}^{2}, \mathrm{H}e^{3} \rangle \mathrm{III}},$$
(19)

depends critically on the relative abundance of He³ to protons. A plot of $F_{\tau}(H^2)$ is shown in Figure 4 for the case where III is assumed equal to its equilibrium value. The range of validity of the assumption of He³ equilibrium is discussed in Section IVa. From equations (14a) and (14b) it is clear that, in the normal situation where the He³ abundance is increasing, the maximum value of the He³-to-proton ratio is obtained from the equilibrium condition. Hence, Figure 4 represents an upper limit on the fraction of deuterium burned by the H²(τ , p)He⁴ reaction in the interiors of stars.



FIG. 4.—The fraction of deuterium consumed by the $H^2(\tau, p)He^4$ reaction is shown as a function of T_6 for $X_H = X_{H_e} = 0.5$. The dashed curve has the same significance as in Fig. 2.

IV. He³-BURNING REACTIONS

A large number of reactions play important roles in the conversion of He³ to He⁴. In the discussion below, we examine the conditions under which each He³-burning reaction plays a significant role.

a) The Abundance of He^3

In all the calculations described in Sections III, IV, and V we have assumed that the abundance of He³ is equal to its equilibrium value (Fig. 2). However, at temperatures below 6×10^6 ° K, the mean lifetime of a He³ nucleus is sufficiently long ($\geq 10^{10}$ years) that it is unlikely that He³ equilibrium can be established during the time that a star of one solar mass spends on the main sequence ($\sim 10^{10}$ years). Hence at temperatures below 6×10^6 ° K the He³ abundance will almost certainly be less than its equilibrium value, perhaps several orders of magnitude less.

For temperatures below $T_6 = 6$, the actual He³ abundance depends on the mass, age, and chemical composition of the star as well as the local temperature. Thus, the He³ abundance cannot be determined in general but must be calculated separately for each case. Lack of He³ equilibrium seriously affects many of the quantities of interest in the proton-proton chain. In this paper, the curves which depend sensitively on the He³ abundance have been calculated for an equilibrium He³ abundance but have been dashed in the temperature region below $T_6 = 6$. We have assumed an equilibrium abundance of He³ in order to call attention to the fact that at very low temperatures (<4 × 10⁶ ° K) it is possible to build up large relative abundances of He³ and also because of the large range of He³ abundances that are possible.

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A lack of He³ equilibrium will have only a small effect on our conclusions regarding the burning of deuterium. The importance of the H²(τ , p)He⁴ reaction will be drastically reduced at low temperatures so that $F_{\tau}(H^2)$ will probably reach a maximum value of about 10⁻³ in the neighborhood of $T_6 = 5$ and then fall rapidly at still lower temperatures. One further result of this decrease in the already minor role of the H²(τ , p)He⁴ reaction as a H²-burning reaction will be a slight increase in the abundance of deuterium at low temperatures. This latter effect, however, is small, amounting to only 5 per cent at $T_6 = 1$ and falling to less than $\frac{1}{3}$ per cent by $T_6 = 5$.

In our discussions of He³ burning, we note the effect of a non-equilibrium abundance of He³ on the various quantities derived. The principal result is a reduction in the importance of the He³(τ , 2p)He⁴ termination at temperatures below $T_6 = 6$.

b) $He^{3}(p, \gamma)Li^{4}$

The importance of the stability, or instability, of Li^4 to proton emission has long been recognized in relation to such questions as the speed and efficiency of the termination of the proton-proton chain, the ages of astronomical systems, and the production of high-energy solar neutrinos (cf. Bethe 1939; Reeves 1959; and Fowler 1960).

Because of the great astrophysical importance of knowing whether Li⁴ is particle stable, Bashkin, Kavanagh, and Parker (1959) investigated the particle stability of this isotope by searching for high-energy positrons from the beta-decay of Li⁴. Bashkin *et al.* established an experimental upper limit on the cross-section for the formation of Li⁴ by the reaction He³(p, γ)Li⁴ that is 25 times smaller than the minimum theoretical crosssection predicted by Christy (1959) if Li⁴ were particle stable and had a 1⁻ or 2⁻ ground state. On this basis Bashkin *et al.* were able to conclude that Li⁴ is probably unstable to proton emission.

Contrary to what has been assumed in previous review articles, we assert that the instability of Li⁴ to proton emission does not completely rule out its possible importance as a termination for the proton-proton chain via the reactions

$\operatorname{He}^{3}(p, \gamma)\operatorname{Li}^{4}(\beta^{+}\nu)\operatorname{He}^{4}$.

If the proton-unstable ground state of Li^4 corresponds to a narrow low-lying resonance in the He³ + p system, then one can show, either by assuming the Breit-Wigner form for the cross-section or by statistical-mechanics arguments, that in an environment of He³'s and protons the equilibrium abundance of Li⁴ is given by

$$n(\mathrm{Li}^{4}) = n(\mathrm{He}^{3}) n(\mathrm{H}^{1}) \omega \left(\frac{8\pi\hbar^{2}}{3m_{p}kT}\right)^{3/2} e^{-E_{r}/kT} \mathrm{cm}^{-3}, \qquad (20)$$

where E_r is the amount by which the Li⁴ ground state is unbound, ω is the statistical weight, $(2J_{Li^4} + 1)/[(2J_{He^3} + 1)(2J_p + 1)] = \frac{1}{4}(2J_{Li^4} + 1)$, and all the other symbols have their usual meaning.

The mean life of He³ for conversion into He⁴ via the Li⁴ termination is

$$\tau_p(\text{He}^3) = \frac{n(\text{He}^3)}{n(\text{Li}^4)} \tau_{\beta^+}(\text{Li}^4), \qquad (21)$$

or, from equation (20),

$$\tau_{p}(\text{He}^{3}) = 0.8205 \times 10^{6} \left[\frac{\tau_{\beta^{+}}(\text{Li}^{4})}{\rho X_{\text{H}}(2J_{\text{Li}^{4}}+1)} \right] T_{6}^{3/2} \exp \frac{E_{r}(\text{keV})}{0.0862T_{6}},$$
(22)

where $\tau_{\beta^+}(\text{Li}^4)$ is the mean Li⁴ positron lifetime. On the basis of theoretical discussions by Breit and McIntosh (1951) and by Werntz and Brennan (1963), we conclude that the mean beta-decay lifetime of Li⁴ lies within the limits

$$10^{-3} \sec \le \tau_{\beta^+}(\text{Li}^4) \le 4 \times 10^3 \sec .$$

This large range in the possible lifetimes is due to the unknown spin and parity of the Li^4 ground state. Shell-model considerations suggest that the ground state may be 1⁻ or 2⁻, and more complicated arguments indicate it may even be 0⁺ (Werntz and Brennan 1963).

Since the properties of the ground state of Li⁴ are at present completely unknown, we have calculated $\tau_p(\text{He}^3)$ for a number of values of E_r as a function of T_6 using equation (22) and assuming $J^{\pi}(\text{Li}^4) = 1^-$ (our results depend only slightly on this assumption) and $\tau_{\beta^+}(\text{Li}^4) = 10^{-3}$, 1.0, and 10^{+3} sec. The results of these calculations are plotted in Figure 5 for $\tau_{\beta^+}(\text{Li}^4) = 1.0$ sec, and $\tau_p(\text{He}^3)$ is compared to $\tau_p(\text{H}^1)$ and $\tau(\text{He}^3)$, the mean



FIG. 5 —The mean life of He³ to the Li⁴ termination is shown as a function of T_6 for $X_H = 0.5$, $\rho = 100$ gm cm⁻³, $J^{\pi}(\text{Li}^4) = 1^-$, $\tau_{\beta^+}(\text{Li}^4) = 1^-$ osc and various values of E_r . For comparison, the mean life of the H¹($\rho, \beta^+\nu$)H² reaction and τ (He³), the mean life of He³ to burning by H², He³, and He⁴, are also shown

life of He³ to the He³-burning reactions (13d), (13e), and (13f). The quantity τ (He³) is defined by

$$\tau(\mathrm{He}^{3})^{-1} = \tau_{d}(\mathrm{He}^{3})^{-1} + \tau_{\tau}(\mathrm{He}^{3})^{-1} + \tau_{\alpha}(\mathrm{He}^{3})^{-1} .$$
⁽²³⁾

Since $\tau_p(\text{He}^3)$ is proportional to $\tau_{\beta+}(\text{Li}^4)$, the curves in Figure 5 can easily be scaled to values of $\tau_{\beta^+}(\text{Li}^4)$ other than 1.0 sec.

The fraction of the He³ consumed that is converted to He⁴ via Li⁴ is

$$F_{p}(\text{He}^{3}) = \frac{\tau(\text{He}^{3})}{\tau(\text{He}^{3}) + \tau_{p}(\text{He}^{3})}.$$
(24)

The quantity $F_p(\text{He}^3)$ is plotted as a function of E_r in Figure 6 for $T_6 = 15$ and for the cases of $\tau_{\beta^+}(\text{Li}^4) = 10^{-3}$, 1.0, and 10^{+3} sec. A comparison with results at other temperatures indicates that, in the temperature region around $15 \times 10^6 \,^{\circ}$ K, $F_p(\text{He}^3)$ is fairly insensitive to temperature A temperature change of $5 \times 10^6 \,^{\circ}$ K shifts the curves in Figure 6 by only 1 or 2 keV.

From Figures 5 and 6 we conclude that E, must be less than 40 keV in order for the

Li⁴ termination to be of importance in the astrophysically interesting region, $5 \le T_6 \le 30$.

Since the completion of the proton-proton chain by the He³ (p, γ) Li⁴ $(\beta^+\nu)$ He⁴ termination would produce a flux of high-energy solar neutrinos, we can use the measurements of Davis (1964) of the solar-neutrino flux to establish a lower limit on the possible value of E_r . Our argument is an elaboration of one described briefly by Bahcall (1964).

Davis (1964) has established an upper limit on the solar-neutrino flux at the earth's orbit of

$$[\varphi_{\nu}\bar{\sigma}_{\nu}(\mathrm{Cl}^{37})] \leq 3 \times 10^{-34} \operatorname{sec}^{-1} \mathrm{Cl}^{37} - \operatorname{atom}^{-1}.$$
(25a)

Bahcall (1964) has shown that the average cross-section for absorption of Li⁴ neutrinos by Cl³⁷ (including transitions to excited states) is



 $\bar{\sigma}_{\nu({\rm Li}^4)}({\rm Cl}^{37}) \simeq 2 \times 10^{-42} \,{\rm cm}^2 \,.$ (25b)

FIG. 6.—The fraction of He³ consumed by the Li⁴ termination is shown as a function of E_r for $X_{\rm H} = 0.5$, $\rho = 100$ gm cm⁻³, $T_6 = 15$, $J^{\pi}({\rm Li}^4) = 1^-$ and $\tau \beta^+({\rm Li}^4) = 10^{-3}$, 1.0, 10³ sec.

This cross-section, combined with the experimental upper limit of Davis, implies the following upper limit on the flux of Li^4 neutrinos at the earth's orbit:

$$\varphi_{\nu}(\text{Li}^4) \le 1.5 \times 10^{+8} \text{ cm}^{-2} \text{ sec}^{-1}$$
 (25c)

Thus the energy flux, $\epsilon(\text{Li}^4)$, at the earth's orbit from proton-proton reactions terminated by Li⁴ in the Sun is less than 2.5 × 10⁺⁹ MeV cm⁻² sec⁻¹. Comparing this energy flux with the solar constant, $\epsilon(\bigcirc) = 8.7 \times 10^{+11}$ MeV cm⁻² sec⁻¹, we find that

$$\frac{\epsilon(\mathrm{Li}^4)}{\epsilon(\odot)} \le 0.29 \times 10^{-2}.$$
(26a)

The fractional energy flux due to Li⁴ terminations can also be expressed as

$$\frac{\epsilon(\mathrm{Li}^4)}{\epsilon(\odot)} = \frac{(16.8)F_p(\mathrm{He}^3)}{(16.8)F_p(\mathrm{He}^3) + (13.1)F_{\mathrm{He}^3}(\mathrm{He}^3) + (25.7)F_{\mathrm{He}^4}(\mathrm{He}^3)}.$$
 (26b)

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It follows from the work of Sears (1964) or Bahcall *et al.* (1963) that $F_{\text{He}^3}(\text{He}^3) \cong 4F_{\text{He}^4}(\text{He}^3)$. Hence, if we neglect the small contribution due to Li⁴, $F_{\text{He}^3}(\text{He}^3) = 0.8$ and $F_{\text{He}^4}(\text{He}^3) = 0.2$. Inserting these numbers in equations (26*a*) and (26*b*), we find

$$F_p(\text{He}^3) |_{\odot} \le 0.27 \times 10^{-2}$$
 (27)

Since

$$C_{\mathrm{He}^4}(\mathrm{He}^3, p) = [1 \times C_{\mathrm{He}^4}(\mathrm{He}^3, \mathrm{He}^4) + 2 \times C_{\mathrm{He}^4}(\mathrm{He}^3, \mathrm{He}^3)] \times F_p(\mathrm{He}^3)$$

we find

$$C_{\text{He}^4}(\text{He}^3, \text{He}^4) \simeq \frac{1}{3}, C_{\text{He}^4}(\text{He}^3, \text{He}^3) \simeq \frac{2}{3}, \text{ and } C_{\text{He}^4}(\text{He}^3, p) |_{\odot} \le 0.45 \times 10^{-2}.$$
 (28)

Finally, comparing this limit on $F_p(\text{He}^3)|_{\odot}$ with Figure 6, we see that the Bahcall (1964), Davis (1964), and Sears (1964) results imply that

$$E_r \ge 20 \text{ keV}$$
, (29)

unless the Li⁴ mean life to beta decay is considerably longer than 4000 sec.

It should be emphasized that at present nothing is known experimentally about the ground state of Li⁴ except its probable instability to proton emission. Hence, although it is a priori unlikely that the Li⁴ ground state lies in the narrow region from 20 to 40 keV above the He³ + p threshold, the only experimental limit that can be placed at present on the astrophysical importance of the Li⁴ termination derives from the solar neutrino measurements of Davis.

b') $He^{3}(p, \beta^{+}\nu)He^{4}$

Even if the ground state of Li⁴ proves to be unbound by several MeV, He³ can be converted into He⁴ by the direct beta-decay reaction He³(p, $\beta^+\nu$)He⁴.

Salpeter (1952) has shown that the He³ $(p, \beta^+\nu)$ He⁴ termination is unimportant for temperatures between $T_6 = 5$ and $T_6 = 30$. An extrapolation of Salpeter's calculations suggests, however, that at temperatures for which $T_6 \leq 1$ the He³ $(p, \beta^+\nu)$ He⁴ reaction may be the fastest He³-burning reaction except for the special conditions discussed in Section IVf. This extrapolation provides, however, only a rough estimate, and a further investigation of the He³ $(p, \beta^+\nu)$ He⁴ reaction is necessary for a full understanding of the operation of the proton-proton chain at temperatures below $T_6 = 1$. At a temperature of 1×10^6 ° K, however, the mean life of He³ to the He³ $(p, \beta^+\nu)$ He⁴ reaction is probably $\sim 10^{22}$ years, and this termination is therefore incapable of producing a He³ equilibrium abundance or of generating any significant amount of energy.

c) $He^{3}(d, p)He^{4}$

This reaction was discussed in Section IIIc in connection with H² burning; we now examine its role as a He³-burning reaction. The fraction of the He³ destroyed that is consumed by the He³(d, p)He⁴ reaction is

$$F_d(\mathrm{He}^3) = \frac{\langle \mathrm{He}^3, \mathrm{H}^2 \rangle \mathrm{II}}{\langle \mathrm{He}^3, \mathrm{H}^2 \rangle \mathrm{II} + \langle \mathrm{He}^3, \mathrm{He}^3 \rangle \mathrm{III} + \langle \mathrm{He}^3, \mathrm{He}^4 \rangle \mathrm{IV}},$$
(30)

and the fraction of the He³ produced that is consumed by the He³(d, p)He⁴ reaction is

$$\Phi_d(\mathrm{He}^3) = \frac{\langle \mathrm{He}^3, \mathrm{H}^2 \rangle \mathrm{III}}{\langle \mathrm{H}^2, \mathrm{H}^1 \rangle}.$$
(31)

The fraction of He⁴ that is produced by the He³(d, p)He⁴ termination is

$$C_{\mathrm{He}^{4}}(\mathrm{He}^{3}, \mathrm{H}^{2}) = \frac{\langle \mathrm{He}^{3}, \mathrm{H}^{2} \rangle \mathrm{II}}{\langle \mathrm{He}^{3}, \mathrm{H}^{2} \rangle \mathrm{II} + \frac{1}{2} \langle \mathrm{He}^{3}, \mathrm{He}^{3} \rangle \mathrm{III} + \langle \mathrm{He}^{3}, \mathrm{He}^{4} \rangle \mathrm{IV}}.$$
(32)

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The quantities $F_d(\text{He}^3)$ and $C_{\text{He}^4}(\text{He}^3, \text{H}^2)$ have been calculated as a function of temperature, assuming an equilibrium abundance of He³; the results are shown in Figures 7 and 8. (The assumption of an equilibrium He³ abundance implies that $\Phi_d[\text{He}^3] = F_d[\text{He}^3]$.)

Figures 7 and 8 show that the He³(d, p)He⁴ reaction is important as a He³-burning reaction only for temperatures below $T_6 = 6$, where we have shown that the He³ abundance is usually less than its equilibrium value. In such a non-equilibrium situation, equations (30), (31), and (32) show that, although the fraction Φ_d (He³) is reduced (not all the He³ produced will be destroyed), both F_d (He³) and C_{He^4} (He³, H²) will be enhanced. Hence at low temperatures ($T_6 \leq 3$), it is likely that whatever He³ is converted into He⁴ is converted via the He³(d, p)He⁴ reaction, except under the special conditions discussed in Sections IV f and b'.



FIG. 7.—The fractions of He³ consumed by the various He³-burning reactions are shown as a function of T_6 for $X_{\rm H} = X_{\rm H_0} = 0.5$ and $\rho = 100$ gm cm⁻³. The He³-consumption fractions were calculated assuming an equilibrium abundance of He³ and neglecting possible Li⁴ terminations. The dashed curves have the same significance as in Fig. 2.

Despite the "relative" importance of the He³(d, p)He⁴ reaction at low temperatures, it should be noted that for $T_6 \leq 3$, τ_d (He³) $\geq 10^{16}$ years and hence this reaction never contributes significantly to the energy generation.

d) $He^{3}(\tau, 2p)He^{4}$

The cross-section for the He³(τ , 2p)He⁴ reaction has been measured by Good, Kunz, and Moak (1954) over a range of He³ energies from 100 to 800 keV. Their low-energy results are not inconsistent with the following intercept parameters for S(E):

$$S_0 \sim 1100 \text{ keV-barns}$$
,

$$(dS/dE)_0 \sim 0$$
 barns.

However, a plot of S versus E_{CM} for the He³ $(\tau, 2p)$ He⁴ reaction using the data of Good *et al.* suggests that the value of S_0 quoted in the literature for this reaction $[S_0 \approx 1100 \text{ keV-barns}]$ may have been determined by the limitations of their experiment, for ex-

ample, by the uncertainty in the average beam energy and by the straggle of the beam through their target. The correct value of S_0 may be as much as a factor of 5 or even 10 different from the value quoted above. Further experimental clarification of this problem is urgently needed.

We have calculated the relevant fractions, $F_{\tau}(\text{He}^3)$, $\Phi_{\tau}(\text{He}^3)$ and $C_{\text{He}^4}(\text{He}^3, \text{He}^3)$, and these are plotted as functions of the temperature in Figures 7 and 8 for the case $X_{\text{H}} = X_{\text{He}} = 0.50$ and $n(\text{He}^3) = 0$. The quantity $C_{\text{He}^4}(\text{He}^3, \text{He}^3)$ is defined by analogy with equation (32)

$$C_{\mathrm{He}^{4}}(\mathrm{He}^{3}, \mathrm{He}^{3}) = \frac{\frac{1}{2} \langle \mathrm{He}^{3}, \mathrm{He}^{3} \rangle \mathrm{III}}{\langle \mathrm{He}^{3}, \mathrm{H}^{2} \rangle \mathrm{II} + \frac{1}{2} \langle \mathrm{He}^{3}, \mathrm{He}^{3} \rangle \mathrm{III} + \langle \mathrm{He}^{3}, \mathrm{He}^{4} \rangle \mathrm{IV}}.$$
(33)



FIG. 8.—The fractions of He⁴ produced by various terminations are shown as a function of T_6 for $X_{\rm H} = X_{\rm H_0} = 0.5$ and $\rho = 100$ gm cm⁻³. The He⁴-production fractions were calculated assuming an equilibrium abundance for He³ and neglecting possible Li⁴ terminations. The dashed curves have the same significance as in Fig. 2.

Since a non-equilibrium abundance of He³ will reduce all three of the fractions, F_{τ} (He³), Φ_{τ} (He³), and C_{He^4} (He³, He³), at temperatures below $T_6 = 6$ the curves in Figures 7 and 8 represent upper limits.

e) $He^{3}(a, \gamma)Be^{7}$

Experimental measurements on the He³(a, γ)Be⁷ reaction by Holmgren and Johnston (1959) and by Parker and Kavanagh (1963) have shown that the cross-section for this reaction is sufficiently large so that at high temperatures the burning of He³ by alphaparticles competes successfully with the He³($\tau, 2p$)He⁴ reaction. From their measurements, Parker and Kavanagh have obtained the following intercept parameters for S(E):

$$S_0 = (0.47 \pm 0.05) \text{ keV-barns},$$

$$(dS/dE)_0 = -2.8 \times 10^{-4}$$
 barns.

The fractions, $F_{\alpha}(\text{He}^3)$, $\Phi_{\alpha}(\text{He}^3)$, and $C_{\text{He}^4}(\text{He}^3, \text{He}^4)$ have been calculated for the $\text{He}^3(\alpha, \gamma)\text{Be}^7$ reaction and are plotted in Figures 7 and 8 for an equilibrium abundance of He³. The quantity $C_{\text{He}^4}(\text{He}^3, \text{He}^4)$ is defined by analogy with equation (32) as

$$C_{\mathrm{He}^{4}}(\mathrm{He}^{3},\mathrm{He}^{4}) = \frac{\langle \mathrm{He}^{3}\,\mathrm{He}^{4}\rangle\mathrm{IV}}{\langle \mathrm{He}^{3},\mathrm{H}^{2}\rangle\mathrm{II} + \frac{1}{2}\langle \mathrm{He}^{3},\mathrm{He}^{3}\rangle\mathrm{III} + \langle \mathrm{He}^{3},\mathrm{He}^{4}\rangle\mathrm{IV}}.$$
⁽³⁴⁾

If the He³ abundance is less than its equilibrium value, then $\Phi_{\alpha}(\text{He}^3)$ is decreased, and both $F_{\alpha}(\text{He}^3)$ and $C_{\text{He}^4}(\text{He}^3, \text{He}^4)$ are increased relative to the values plotted in Figures 7 and 8. However, at the low temperatures where the non-equilibrium abundance of He³ is important, the He³(α , γ)Be⁷ reaction is so infrequent [$F_{\alpha}(\text{He}^3) \leq 10^{-4}$ at $T_6 = 6$; $\leq 10^{-9}$ at $T_6 = 2$], that we can safely ignore non-equilibrium considerations for this reaction.

The following reactions are possible ways to convert Be⁷ into two He⁴'s and terminate the proton-proton chain:

(a) Be⁷(e⁻, ν)Li⁷(p, a)He⁴;
(b) Be⁷(p, γ)B⁸(β⁺ν)Be^{8*}(a)He⁴;
(c) Be⁷(d, p)Be⁸(a)He⁴;
(d) Be⁷(τ, p)B⁹(p)Be⁸(a)He⁴, Be⁷(τ, a)Be⁶(p)Li⁵(p)He⁴; and
(e) Be⁷(a, γ)C¹¹(β⁺ν)B¹¹(p, a)Be⁸(a)He⁴.

Of these reactions, only (a) and (b) are sufficiently fast to guarantee equilibrium; (c) is too slow because of the small relative abundance of deuterium and (d) and (e) are too slow because of their larger coulomb barriers.

From Figure 3, *B*, we can see that the $\text{Li}^7(p, a)\text{He}^4$ reaction is sufficiently fast at all temperatures of interest so that the rate of equation (35*a*) is determined by the Be⁷(e^- , ν)Li⁷ reaction. The positron mean life of B⁸ in stellar interiors does not differ significantly (Bahcall 1962*a*) from its terrestrial value of about 1 sec, and so from Figure 3, *B*, we can see that the rate of equation (35*b*) is determined by the Be⁷(p, γ)B⁸ reaction.

Bahcall (1962b) and Kavanagh (1960) have studied terminations (35a) and (35b). Bahcall has shown that for the $Be^7(e^-, \nu)Li^7$ reaction the mean lifetime of Be^7 for continuum electron capture in stars like the Sun is

$$\tau_{e^{-}}(\mathrm{Be}^{7}) = \frac{10^{9} T_{6}^{1/2}}{2.12\rho \left(1 + X_{\mathrm{H}}\right)} \sec.$$
(36)

From Kavanagh's measurements of the cross-section for the $Be^7(p, \gamma)B^8$ reaction at proton energies of 800 and 1400 keV, Christy and Duck (1961) have obtained the following intercept parameters:

$$S_0 = (0.030 \pm 0.010)$$
 keV-barns, $(dS/dE)_0 = 0$ barns.

The fraction of Be^7 consumed via reaction (35*a*) is

$$F_{e^-}(\mathrm{Be}^7) = \frac{\tau_p(\mathrm{Be}^7)}{\tau_{e^-}(\mathrm{Be}^7) + \tau_p(\mathrm{Be}^7)},$$
(37a)

and the fraction consumed via reaction (35b) is

$$F_{p}(\text{Be}^{7}) = \frac{\tau_{e^{-}}(\text{Be}^{7})}{\tau_{e^{-}}(\text{Be}^{7}) + \tau_{p}(\text{Be}^{7})}.$$
(37b)

From equations (37a) and (37b) we find that the fraction of He⁴ produced through each of these terminations is

$$C_{\rm He^4}({\rm Be^7}, e^-) = C_{\rm He^4}({\rm He^3}, {\rm He^4})F_{e^-}({\rm Be^7}), \qquad (38a)$$

$$C_{\text{He}^4}(\text{Be}^7, \text{H}^1) = C_{\text{He}^4}(\text{He}^3, \text{He}^4)F_p(\text{Be}^7)$$
 (38b)

The two fractions, $C_{\text{He}^4}(\text{Be}^7, e^-)$ and $C_{\text{He}^4}(\text{Be}^7, \text{H}^1)$, are plotted in Figure 8 as functions of temperature and are compared with the C_{He^4} 's for the terminations discussed in Sections IVc and IVd.

f) $He^{4}(e^{-}, \nu)T^{3}$

At low temperatures the He³-burning reactions (13d), (13e), and (13f) occur more slowly than the He³-producing reactions (13a) and (13b). Hence at such temperatures the proton-proton chain would effectively end at He³ if the only reactions possible were those listed in equation (13). Schatzman (1958) has, however, discussed an alternative way of closing the proton-proton chain via the reactions

$$He^{3}(e^{-}, \nu)H^{3}$$
, (39a)

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$$\mathrm{H}^{3}(p, \gamma)\mathrm{He}^{4}$$
. (39b)

Bahcall and Wolf (1964) have recently reinvestigated terminations (39a) and (39b) using the modern theory of nuclear beta-decay and have also considered some alternative ways of burning the tritium produced by reaction (39a). The reader is referred to their accompanying paper for further details. In our discussion we summarize some of the conclusions of Bahcall and Wolf. Iben (1963) is currently studying models for stars of less than 1 solar mass in order to determine under what conditions reactions (39a) and (39b) actually terminate the proton-proton chain.

The reaction $\operatorname{He}^{3}(e^{-}, \nu)\operatorname{H}^{3}$ is an endothermic electron-capture reaction with a threshold energy of about 18 keV (Langer and Moffat 1952). Bahcall and Wolf have shown that the rate of the $\operatorname{He}^{3}(e^{-}, \nu)\operatorname{H}^{3}$ reaction is negligibly small for densities less than a critical density,

$$\rho_{\rm cr} = 1.87 \mu_e \times 10^{+4} \text{ gm cm}^{-3}$$
,

where μ_{e} is the mean molecular weight per electron. This critical density is determined by the requirement that the Fermi energy for a completely degenerate gas of electrons be equal to the threshold energy of 18 keV.

The electron-capture lifetime for a He³ nucleus is

$$\tau_{1/2}(\text{He}^3) = (\text{ft}_{1/2})_{\text{lab}} K^{-1} , \qquad (40)$$

where $(ft_{1/2})_{lab}$ is the terrestrial ft-value for the exothermic tritium beta-decay and is equal to 1070 ± 70 sec (Porter 1959) and K is the generalized phase-space factor. Accurate analytic and numerical expressions for K have been given by Bahcall and Wolf for both non-degenerate and degenerate situations. For degenerate situations with densities such that $\rho_{\rm er} < \rho < 10^6$ gm gm⁻³, they find

$$K = 8.22 \times 10^{-10} (E_F - 18)^3 [1 + 0.019 (E_F - 18)].$$
⁽⁴¹⁾

In equation (41) E_F is the Fermi energy in keV, that is,

$$E_F = 511 \left\{ \left[1.018 \left(\frac{\rho}{\mu_e} \right)^{2/3} \times 10^{-4} + 1 \right]^{1/2} - 1 \right\} \text{ keV}.$$
⁽⁴²⁾

Perry and Bame (1955) studied the $H^{3}(p, \gamma)He^{4}$ reaction at center-of-mass energies

from 75 keV to 4.7 MeV, and from their measurements one can obtain the following intercept parameters for the cross-section factor

$$S_0 = 1.5 \times 10^{-3} \text{ keV-barns}$$
,
 $(dS/dE)_q = 2.7 \times 10^{-5} \text{ barns}$.

Two other tritium-burning reactions of interest are $H^{3}(t, 2n)He^{4}$ and $H^{3}(\tau, np)He^{4}$. We estimate on the basis of work by Jarmie and Allen (1958) and Youn, Osetinskii, Sodnom, Govorov, Sizov, and Salatskii (1961) that these reactions have S_{0} 's of 300 keV-barns and 1000 keV-barns, respectively. Using these intercept parameters Bahcall and Wolf have shown that the mean lifetime of tritium to these three nuclear reactions is less than the electron-capture lifetime of He³ at all relevant stellar temperatures and densities. Hence, the lifetimes for the terminations via $He^{3}(e^{-}, \nu)H^{3}$ are determined entirely by the electron-capture lifetime given by equation (40).

The electron-capture lifetime of He³ for the He³(e^- , ν)H³ reaction is only 5000 years for a density of 2.2 × 10⁴ μ_e gm cm⁻³($E_F = 20$ keV). Thus at densities only slightly greater than $\rho_{\rm or}$ the He³(e^- , ν)H³ reaction is considerably faster than the H¹(p, $\beta^+\nu$)H² reaction at all temperatures of interest for main-sequence stars. Hence, at sufficiently high densities reactions (39*a*) and (39*b*) can establish an equilibrium abundance of He³.

g) Solar He³ Burning

In conclusion, we note that stellar-model calculations by Iben and Sears (1962) suggest that over the Sun as a whole about two-thirds of the proton-proton chain terminations occur via the He³(τ , 2p)He⁴ reaction, one-third via the He³(a, γ)Be⁷(e^- , ν)Li⁷ reactions, and 1/1000 via the He³(a, γ)Be⁷(p, γ)B⁸ reactions. The ratio of He³(τ , 2p)He⁴ completions to He³(a, γ)Be⁷ completions is approximately reversed at the center of the Sun compared to the average values quoted above.

v. He⁴ production and energy generation

Over most of the temperature range in which the proton-proton chain operates the $H^1(p, \beta^+\nu)H^2$ reaction is the slowest link in the chain. However, the rate of He⁴ production, R_{α} , is not proportional to the rate of the $H^1(p, \beta^+\nu)H^2$ reaction, $\frac{1}{2}\langle H^1, H^1\rangle n(H^1)n(H^1)$. Instead we must write

$$R_{\alpha} = \chi_{\alpha} [\frac{1}{2} \langle \mathrm{H}^{1}, \mathrm{H}^{1} \rangle n(\mathrm{H}^{1}) n(\mathrm{H}^{1})] \mathrm{cm}^{-3} \mathrm{sec}^{-1}, \qquad (43)$$

where χ_{α} is a dimensionless function of temperature. The temperature variation of χ_{α} arises because the terminations $\operatorname{He}^{3}(d, p)\operatorname{He}^{4}$ and $\operatorname{He}^{3}(\tau, 2p)\operatorname{He}^{4}$ both require two $\operatorname{H}^{1}(p, \beta^{+}\nu)\operatorname{H}^{2}$ reactions for each He^{4} produced $(\chi_{\alpha} = \frac{1}{2})$ while the $\operatorname{He}^{3}(\alpha, \gamma)\operatorname{Be}^{7}$ terminations produce one He^{4} for every $\operatorname{H}^{1}(p, \beta^{+}\nu)\operatorname{H}^{2}$ reaction $(\chi_{\alpha} = 1)$. Writing R_{α} explicitly,

$$R_{\alpha} = \langle \operatorname{He}^{3}, \operatorname{H}^{2} \rangle n(\operatorname{He}^{3})n(\operatorname{H}^{2}) + \frac{1}{2} \langle \operatorname{He}^{3}, \operatorname{He}^{3} \rangle n(\operatorname{He}^{3})n(\operatorname{He}^{3}) + \langle \operatorname{He}^{3}, \operatorname{He}^{4} \rangle n(\operatorname{He}^{3})n(\operatorname{He}^{4}),$$
(44)

we see that

$$\chi_{a} = \frac{\langle \text{He}^{3}, \text{H}^{2} \rangle \text{III II} + \frac{1}{2} \langle \text{He}^{3}, \text{He}^{3} \rangle \text{III}^{2} + \langle \text{He}^{3}, \text{He}^{4} \rangle \text{III IV}}{\frac{1}{2} \langle \text{H}^{1}, \text{H}^{1} \rangle}.$$
 (45)

The quantity χ_{α} is plotted as a function of temperature in Figure 9 assuming that $X_{\rm H} = X_{\rm He} = 0.5$ and that the He³ abundance is equal to its equilibrium value.

At low temperatures, the He³ abundance is less than its equilibrium value and the proton-proton chain tends to terminate with the production of He³. Thus at low temperatures χ_{α} is much less than the equilibrium values shown in Figure 9 and, in fact, ap-

proaches zero as the temperature goes to zero. For the conditions in which the chain is terminated by the He³(e^- , ν)H³(p, γ)He⁴ reactions, χ_{α} is equal to unity. Note that χ_{α} is one-half the quantity φ_{α} defined by Fowler (1958, 1960); the latter quantity was defined in terms of the He³(τ , 2p)He⁴ reaction instead of the H¹(p, $\beta^+\nu$)H² reaction.

The rate of useful energy generation (i.e., energy not carried off by neutrinos) is not proportional to the rate of the $H^1(p, \beta^+\nu)H^2$ reaction or even to R_{α} . If we write the rate of useful energy generation, R_{ϵ} , as

$$R_{\epsilon} = \chi_{\epsilon} [\frac{1}{2} 26.73 \langle \mathrm{H}^{1}, \mathrm{H}^{1} \rangle n(\mathrm{H}^{1}) n(\mathrm{H}^{1})] \mathrm{MeV} \mathrm{cm}^{-3} \mathrm{sec}^{-1}, \qquad (46)$$



FIG. 9.—The He⁴-production and energy-generation factors, χ_a and χ_e , are shown as functions of T_6 for $X_{\rm H} = X_{\rm He} = 0.5$ and $\rho = 100$ gm cm⁻³. An equilibrium abundance is assumed for He³ and any Li⁴ terminations are neglected. The dashed curves have the same significance as in Fig. 2.

where 26.73 MeV is the Q-value for converting $4H^1$ into He^4 , then the additional variation of χ_{ϵ} is caused by the differences in the neutrino-energy losses of the various terminations. The "useful" Q-values for the various terminations of the proton-proton chain are (König, Mattauch, and Wapstra 1962):

	MeV
$\operatorname{He}^{3}(\rho, \beta^{+}\nu)\operatorname{He}^{4}\dots\dots\dots$ 16.8	
$He^{3}(\tilde{d}, p)He^{4} 26.22$	MeV
$He^{3}(\tau, 2p)He^{4}$ 26.22	MeV
$He^{3}(a, \gamma)Be^{7}(e^{-}, \nu)Li^{7}(p, a)He^{4}$ 25.67	MeV
$\mathrm{He}^{3}(a, \gamma)\mathrm{Be}^{7}(p, \gamma)\mathrm{B}^{8}(\beta^{+}\nu)\mathrm{Be}^{8*}(a)\mathrm{He}^{4}$ 19.1	MeV
$\operatorname{He}^{3}(e^{-}, \nu)\operatorname{H}^{3}(\bar{p}, \gamma)\operatorname{He}^{4}\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots\ldots$	MeV

Considering only the burning of He³ by H², He³, and He⁴, we can write R_{ϵ} explicitly as

$$R_{e} = 26.22 \langle \text{He}^{3}, \text{H}^{2} \rangle n(\text{He}^{3})n(\text{H}^{2}) + \frac{1}{2}26.22 \langle \text{He}^{3}, \text{He}^{3} \rangle n(\text{He}^{3})n(\text{He}^{3}) \\ + 25.67 \langle \text{He}^{3}, \text{He}^{4} \rangle F_{e} - (\text{Be}^{7})n(\text{He}^{3})n(\text{He}^{4}) \\ + 19.1 \langle \text{He}^{3}, \text{He}^{4} \rangle F_{p}(\text{Be}^{7})n(\text{He}^{3})n(\text{He}^{4})$$
(47)

It should be noted that with a few refinements χ_{ϵ} is essentially one-half the quantity $\psi(a)$ defined by Fowler (1958, 1960), since the latter quantity was defined in terms of one

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 $He^{3}(\tau, 2p)He^{4}$ reaction instead of one $H^{1}(p, \beta^{+}\nu)H^{2}$ reaction. Thus we can express χ_{ϵ} as

$$\frac{1}{2}\chi_{e} = 0.981 \frac{\langle \text{He}^{3}, \text{He}^{2} \rangle}{\langle \text{H}^{1}, \text{H}^{1} \rangle} \text{III II} + \frac{1}{2}0.981 \frac{\langle \text{He}^{3}, \text{He}^{3} \rangle}{\langle \text{H}^{1}, \text{H}^{1} \rangle} \text{III III} + 0.960 \frac{\langle \text{He}^{3}, \text{He}^{4} \rangle}{\langle \text{H}^{1}, \text{H}^{1} \rangle} F_{e^{-}}(\text{Be}^{7}) \text{III IV} + 0.715 \frac{\langle \text{He}^{3}, \text{He}^{4} \rangle}{\langle \text{H}^{1}, \text{H}^{1} \rangle} F_{p}(\text{Be}^{7}) \text{III IV},$$
or
$$\chi_{\epsilon} = \chi_{\alpha}[0.981C_{\text{He}^{4}}(\text{He}^{3}, \text{H}^{2}) + 0.981C_{\text{He}^{4}}(\text{He}^{3}, \text{He}^{3}) + 0.960C_{\text{He}^{4}}(\text{Be}^{7}, e^{-}) + 0.715C_{\text{He}^{4}}(\text{Be}^{7}, \text{H}^{1})].$$
(48)

The factor χ_{ϵ} is plotted as a function of temperature in Figure 9 assuming an equilibrium He³ abundance and $X_{\rm H} = X_{\rm He} = 0.5$. When the proton-proton chain tends to stop at



FIG. 10.—The energy-generation factor, χ_{ϵ} , is shown as a function of T_6 for $\rho = 100$ gm cm⁻³ and for various values of $(X_{\rm H}, X_{\rm H_0})$. An equilibrium He³ abundance is assumed and Li⁴ terminations are neglected. The dashed curve has the same significance as in Fig. 2.

the production of He³, only 6.68 MeV of useful energy are produced for each H¹(p, $\beta^+\nu$)H² reaction. In such a situation, the He³ abundance is less than its equilibrium value, and χ_{ϵ} approaches a value of 0.250.

The energy generation in the pp-chain can be calculated explicitly from the following expression based on $S_0(pp) = (3.36 \pm 0.4) \times 10^{-22}$ keV-barns:

$$E_{pp} = (4.1 \pm 0.5)\rho X_{\rm H}^2 \chi_{\epsilon} T_6^{-2/3} \exp(-33.804T_6^{-3/2}) \times (1 + 0.012T_6^{1/3} + 0.008T_6^{2/3} + 0.00065T_6 + 0.188\zeta \rho^{1/2}T_6^{-3/2}) \ 10^6 \ {\rm ergs \ gm^{-1} \ scc^{-1}} \,.$$
⁽⁵⁰⁾

All calculations discussed thus far in this paper were made assuming that $X_{\rm H} = X_{\rm He} = 0.5$. Many of the quantities we have calculated depend strongly on $X_{\rm H}$ and $X_{\rm He}$. In order to illustrate this dependence we have plotted in Figure 10 χ_{ϵ} as a function of temperature for hydrogen and helium abundances of $(X_{\rm H} = 0.9; X_{\rm He} = 0.1)$, $(X_{\rm H} = 0.5; X_{\rm He} = 0.5)$, and $(X_{\rm H} = 0.1; X_{\rm He} = 0.9)$.

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