

Direct capture cross sections at low energy

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We derive simple expressions for the low-energy behavior of direct radiative capture cross sections, particularly those important in solar nucleosynthesis, such as ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$, ${}^3\text{H}(\alpha, \gamma){}^7\text{Li}$, and ${}^7\text{Be}(p, \gamma){}^8\text{B}$. Our results for the ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ reaction are inconsistent with a measurement by Rolfs *et al.*

[NUCLEAR REACTIONS ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$, ${}^3\text{H}(\alpha, \gamma){}^7\text{Li}$, $E < 300$ keV, ${}^7\text{Be}(p, \gamma){}^8\text{B}$, $E < 100$ keV, extrapolated S.]

There recently been much activity concerning reevaluation of the cross sections of certain solar nuclear reactions due to the unexpectedly low neutrino flux measured by Davis *et al.*¹ One of these reactions, ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$, is very difficult to measure at solar energies but has a direct bearing on the neutrino problem, since the expected neutrino flux from the sun is almost proportional to the low-energy cross section for this reaction.² Rolfs *et al.*³ have remeasured the low-energy S factor (related to the cross section σ by $S = \sigma E \exp(2\pi\eta)$, where E is the entrance channel center-of-mass energy and η the usual Sommerfeld parameter) and claim it has considerably less energy dependence than was found in previous experiments⁴ or predicted theoretically.⁵ If this were correct, it would substantially alleviate the solar neutrino problem, but would not eliminate it entirely.² Our purpose in this paper is to outline relatively simple expressions for the low energy behavior of direct capture cross sections which can be used to extrapolate the high energy ($E \sim 500$ keV) experimental results for ${}^3\text{He}(\alpha, \gamma){}^7\text{Be}$ to solar energies ($E \sim 20$ keV). We find essential agreement with previous work,^{4,5} and with recent Caltech measurements,⁶ but are in strong disagreement with the data of Ref. 3.

Our treatment is based on the direct capture formalism developed previously,⁷ in which the entrance channel and final state are described by the relative motion of ${}^3\text{He}$ and ${}^4\text{He}$ clusters. Since only $E1$ and $E2$ transitions to the p -wave bound state from s, d and p, f continuum states, respectively, are important at low energy, the S factor for capture to a given final state are

$$S(E1) = k_\gamma^3 (|I_{10}|^2 + |I_{12}|^2), \tag{1}$$

$$S(E2) = \frac{75}{98} k_\gamma^5 (|I_{21}|^2 + \frac{3}{2} |I_{23}|^2),$$

where k is the photon wave number, and

$$I_{\lambda l} = \int_0^\infty r^2 dr \left[\psi_f(r) r^\lambda \frac{\varphi_l(r)}{kr} \right] e^{\pi\eta} (2\pi\eta)^{1/2}. \tag{2}$$

Here, φ_l is the continuum wave function describing the l th partial wave, while ψ_f is the radial p -wave bound-state wave function. (For ${}^7\text{Be}$ the final state has either total spin $j_f = \frac{3}{2}$ with binding energy relative to the entrance channel of 1.586 MeV, or $j_f = \frac{1}{2}$ with binding energy 1.157 MeV.) Note that we have omitted many (common) energy independent factors in Eqs. (1) (including a bound-state spectroscopic factor), as it is only the energy dependence of S which concerns us here.

At low energies, it is well known that the integrand in Eq. (2) peaks at radii far outside the range of the strong interactions.⁷ Therefore, with good accuracy, ψ_f may be replaced with the exponentially decaying $l=1$ Coulomb wave function (related to a Whittaker function). In the same spirit, φ_l may be replaced by a Coulomb wave, modified by the strong interaction phase shift. This we take to correspond to a hard sphere of radius r_0 (see Table I), adjusted to reproduce the elastic scattering,⁵ so that

$$\varphi_l(r) = F_l(kr) - \frac{F_l(kr_0)}{G_l(kr_0)} G_l(kr), \tag{3}$$

where k is the entrance channel wave number and F_l and G_l are the usual regular and irregular Coulomb wave functions. Consistent with this description, we limit the integral in Eq. (2) to $r > r_0$.

To obtain tractable expressions for the $I_{\lambda l}$, it is useful to employ the Bessel function expansions of F_l and G_l (Ref. 8) to generate series in powers of E , each term involving E -independent radial integrals of Bessel functions, powers, and Whittaker functions. These can be readily evaluated numerically by Laguerre quadrature to ob-

TABLE I. Kinematic parameters and results.

	Binding (MeV)	Branching (%)	r_0 (fm)	a (MeV ⁻¹)	b (MeV ⁻²)
³ He(α, γ) ⁷ Be					
$j_f = \frac{3}{2}$	1.586	73	2.8 ^a	-0.575	-0.005
$j_f = \frac{1}{2}$	1.157	27			
³ H(α, γ) ⁷ Li					
$j_f = \frac{3}{2}$	2.468	73	2.8 ^a	-2.034	-3.709
$j_f = \frac{1}{2}$	1.989	27			
⁷ Be(p, γ) ⁸ B	0.136		4.1 ^b	-2.350 ^c	28.3 ^c

^a Reference 5.^b Reference 7.^c Equation (4) valid only for $E \leq 100$ keV.

tain the low-energy behavior of S .

We have considered the reactions ³H(α, γ)⁷Li and ⁷Be(p, γ)⁸B as well as that of primary interest, ³He(α, γ)⁷Be, and have calculated the low energy logarithmic derivative of S in the form

$$\frac{1}{S(E)} \frac{dS}{dE} = a + bE. \quad (4)$$

For the α -capture reactions, where two final states are involved, we have taken the low energy branching ratio to be energy independent and equal to the experimental value.⁹ The kinematic parameters for the three reactions and our results for a and b are shown in Table I. Our value for a for ³He(α, γ)⁷Be agrees with more involved numerical calculations⁵ and is consistent with

previous⁴ and as yet unpublished⁶ Caltech results; it is inconsistent with the data of Ref. 3. Our value for a varies by less than 5% for changes in r_0 of 10%. For ⁷Be(p, γ)⁸B, we find the previously expected⁹ sharp rise in the S factor below $E = 100$ keV; at energies greater than this, the expansion implied by (4) does not converge, due to the small binding energy of the final state.

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