Analytic expression of triple- α reaction rates by a non-adiabatic three-body model

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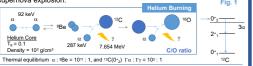
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Triple-α reaction rates have been determined well with the sequential process via the narrow resonances. However, direct triple-α process at off-resonant energies still remains in unsolved problems. In this poster, the direct triple- α process is described with a non-adiabatic Faddeev HHR expansion method. The direct 3α contribution is confirmed to be 10^{-15} _10-3 pb order in photo-disintegration of ${}^{12}\text{C}(2^*$ _1 \rightarrow 0+) for 0.15 < E < 0.35 MeV. Using a three-body type of S-factors, the rates are given in a simple analytic expression. In addition, they are converted into REACLIB format. From composition ratio in NACRE, the non-resonant sequential process between α + bound ${}^{8}\text{Be}$ is found to dominate the NACRE rates for 0.03 < T $_{9}$ < 0.07. I, therefore, find that the current evaluated rates could be reduced by about 10^{-4} at T_{9} = 0.05, from the accurate description of ${}^{8}\text{Be}$ break-up.

1.1 Introduction

Triple-α reaction plays an important role in nucleosynthesis heavier than 12 C, because no stable nuclei exist in mass number A=5 and A=8 [1,2]. This reaction, followed by 12 C(α , γ) 16 D [3], controls C/O ratio at the end of helium burning phase in stars, and it affects up to the nucleosynthesis in e.g. supernova explosion.



I describe direct triple- α process with a non-adiabatic Faddeev HHR*, and I discuss the 3α reaction rates. After I review Faddeev HHR used in the present poster

I deduce a simple analytic form of the rates with introducing S-factors. The rates are also converted into REACLIB. I show that the evaluated rates could be reduced at T₉= 0.05, because of an accurate description of ⁸Be break-up

1.2 Triple-α process

Triple-α reaction via the resonances. (Fig. 1)

In contrast to $^{12}\text{C}(\alpha,\gamma)^{16}\text{C}$, triple- α reaction is currently well-understood through the experimental studies of the 0^{+}_{2} state in ^{12}C (E_{R} = 0.379 MeV).

 E_R is the c.m. energy with respect to the 3α threshold in 12 C. *i.e.* the reaction rates have been determined successfully with the i.e. the reaction rate quential process via the narrow resonances. (e.g. [4,5])

Pioneering work: CF88 [6], Nomoto (1985) [7]

NACRE (1999) [5]

Experimer (1986) [8]. update from CF88, based on Nomoto (1985) [7], Langanke

ned to be bound. The reaction proceeds via two resonances ⁸Be (0+₁), ¹²C (0+₂). Recent experimental progress is found in e.g. [4].

Triple- α reaction from ternary continuum states, (Fig. 2)

- This process is generally expected to be $\it very$ slow, because three $\it \alpha$ -particles almost simultaneously collide and fuse into a 12 C nucleus.
- The direct process is neglected or is treated in some approximations

Adiabatic approximation of 8Be continuum has

1.3 Theoretical studies of triple-α reaction

been applied to tackle the 3α continuum problem Formulae in hyper-spherical coordinates (e.g. [9])

Coulomb Modified Faddeev (CMF) method [10]

Adiabatic channel function (ACF) method [11]

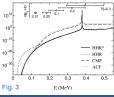
ever, quantum-mechanical description a

off-resonant energies still seems to remain in insolved problen

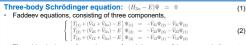
Non-adiabatic approaches of 3α have also been performed, recently. [12,13] (Fig.3)

) lax the continuum states of 8Re

- Faddeev Hyperspherical Harmonics and *R*-matrix (**HHR**) expansion method [13.14.15]. My calculation is labeled with HHR*



2.1 Faddeev HH expansion



Three identical sets of equations are found, because of the symmetric 3a system Third component is rewritten as

Third component is rewritten as
$$\begin{bmatrix} -\frac{\hbar^2}{2g_{12}}\nabla_{2_3}^2 - \frac{\hbar^2}{2g_{1(2)3}}\nabla_{y_3}^2 + \dot{V}_{sym} \end{bmatrix} \psi_{lm} = E \, \psi_{lm} \quad \dot{V}_{sym} = \sum_{i < j}^3 V_{sm}(r_{ij}) + V_{so} \quad \textbf{(3)}$$
 Coupled-channel (CC) equations with hyper-radial wavefunctions:

 $\left[T_{\gamma} + U_{\gamma\gamma}^{l}(\rho) - \epsilon\right]\chi_{\gamma}^{l}(\rho) = -\sum_{l}U_{\gamma'\gamma}^{l}(\rho)\chi_{\gamma'}^{l}(\rho)$ Translate Jacobi coordinates into hyper-spherical coordinates $T_{\gamma} = \frac{d^2}{da^2} - \frac{(K + 3/2)(K + 5/2)}{a^2}$ y_i y_i $\theta_i = \arctan\left(\frac{x_i}{y_i}\right)$ (4) $U^l_{\gamma'\gamma} = -\frac{2m_N}{\hbar^2}V^l_{\gamma'\gamma}(\rho),$ (5) The coordinates are different from CMF

Introducing hyper-angular momentum K, I obtain the *ordinary* CC equations for inelastic scattering. e.g. [16,17] (if L=K+3/2).



The final results are independent of the adopted $\varphi_n^K(\rho)$, if a large number of basis functions are used so as to expand well the wave functions.

2.2 R-matrix expansion

CC equations of Eq.(5): $(T+U)X = \epsilon X$ $E(l_i^+) = -\frac{\hbar^2}{2m_N}\epsilon_i$ $\chi^l_{\gamma i}(\rho) = \sum_n c^n_{\gamma i} \varphi^K_n(\rho)$ Sound states, $\chi^l_{\gamma i}(\rho) \rightarrow \mathcal{N}^l_{\gamma i} W_{-\eta_{\gamma},K+2}(2\kappa\rho)$ $\rightarrow \mathcal{N}_{\gamma i}^{l}W_{-\eta_{\gamma},K}$ (9)

 $E(I^*_i)$ < 0, Bound states, $\chi^l_{\gamma i}(\rho) \to \chi^l_{\gamma i} W_{-\eta_{\gamma},K+2}(2\kappa\rho)$ N: normalization constants, W: Whitterker functions

E(h) > 0, Resonances corresponding to low-lying levels & mathematically orthogonal states without specific interpretation in physics.



S-matrix is defined by R-matrix, Incoming Coulomb function Outgoing Coulomb $S_{\alpha\beta}^{l}(E, a_c) = [Z_{\alpha\beta}^{*}(E, a_c)]^{-1}Z_{\alpha\beta}(E, a_c),$ $Z_{\alpha\beta}(E, a_c) = H_{K+3/2}^{-1}(\eta_0; ka_c)\delta_{\alpha\beta} - a_cR_{\alpha\beta}^{l}(E, a_c)H_{K+3/2}^{-1}(\eta_S; ka_c)$ (12) $\begin{array}{|c|c|} \hline \textbf{\textit{R-matrix}} & R^{l}_{\alpha\beta}(E,a_{c}) & = & \sum_{i} \frac{\tilde{\gamma}_{\alpha i}\tilde{\gamma}_{\beta i}}{E(l^{+}_{i})-E} \end{array}$ (14) Reduced width amplitudes are defined as $\tilde{\gamma}_{ai} = \sqrt{\frac{\hbar^2}{2m_N a_c}} \chi_{ai}^{\rm f}(a_c)$

solved numerically from $\rho = a_c$ to $\rho = \rho_m$, $\chi_{\alpha\beta}^{l,jn}(k,a_c) \longrightarrow \chi_{\alpha\beta}^{l,ext}(k,\rho)$. I use R-matrix propagation technique [13] to obtain the linearly-independent solutions.

Expansion by the linearly-independent waves:

 $\tilde{\chi}_{\gamma\gamma_0}^l(k, \rho) = \sum_l C_{\gamma'\gamma_0}(k) \chi_{\gamma\gamma'}^{l \ ext}(k, \rho)$

. The coefficients are obtained by matching to the asymptotic form of w.f. [12],

 $\tilde{\chi}_{\gamma\gamma_0}^l(k, \rho) \rightarrow \frac{i}{2} \left[I_{\gamma, K+3/2}^{(\gamma_0)}(\eta_{\gamma}; k\rho_m) - \sum_i S_{\gamma'\gamma_0}^l(E) O_{\gamma, K+3/2}^{(\gamma')}(\eta_{\gamma}; k\rho_m) \right]$

Coupled-Coulomb waves [17]: $O_{\gamma,K+3/2}^{(\gamma')}(\eta_{\gamma};k\rho) = a_{\gamma}^{(\gamma')}(k,\rho)H_{\gamma,K+3/2}^{+}(\eta_{\gamma};k\rho), I = O^{*}$ (18) screening, $a_{\gamma}^{(\gamma')} \rightarrow \delta_{\gamma\gamma'}$

+ screening, $a_{\gamma}^{(\gamma)} \rightarrow \delta_{\gamma\gamma'}$ Effective, if the off-diagonal part of Coulomb coupling potentials is relatively small at a matching radius p_{m} , compared with E. (e.g.[16]) Interior scattering waves including long-range Coulomb couplings are obtained as



B(E2) strength between 0* continuum states and 2*, are calculated with Eqs. (8) (16) and (19). Quadruple precision is required to execute stable calculations.

$$B(E2) \ \, \text{strength:} \ \, \frac{dB(E2; \alpha^* \to 2_1^*)}{dE} = \frac{5e^2}{2\pi\hbar i c} \sum_{n} |\sum_{\gamma_1} \int_{\mathcal{K}_{\gamma_1}^{p,q_1}(c)} (\sum_{\gamma_1, \gamma_2} f_{\gamma_1}^{p,q_2}(c) + \int_{\gamma_2, \gamma_2}^{p,q_2} f_{\gamma_2}^{p,q_2}(c) \right) \\ = \frac{[\text{Hyper-radial part}]}{[\text{Hyper-radial part}]} - M_{\sigma_1}^{p,q_2}(c) = M_{\sigma_1}^{p,q_2}(c) + M_{\sigma_2}^{p,q_1}(c) + M_{\sigma_1}^{p,q_2}(c) + H_{\sigma_2}^{p,q_2}(c) + H_{\sigma_2}$$

 $M_{\gamma'\gamma\gamma_0}^{2^+_10^+Ext}(k) = i^K \int_{a_-}^{\infty} \chi_{\gamma'1}^{2^+}(\rho) \tilde{\chi}_{\gamma\gamma_0}^{0^+}(k, \rho) \rho^2 d\rho$ (23)

Photo-disintegration cross sections:
$$\sigma_g(E) = \frac{4\pi^3}{75} \left(\frac{E_g}{hc}\right)^3 \frac{1}{5} \frac{dB(E2;0^+ \to 2_1^+)}{dE}$$
 (24)
Reaction rates: $R_{50}(E) = N_A^2 \frac{489\pi}{dB(24000\pi)^{3/2} c^2} \frac{h^3}{E^2} F_g(E)$ (25)

 $\begin{array}{ll} \dots & \dots & \frac{A}{(\mu_{12}\mu_{(12)3})^{3/2}}\frac{z^2}{c^2}\frac{z^2}{E^2}\sigma_g(E) \\ \langle R_{3\alpha} \rangle & = & \frac{1}{2}\frac{1}{(k_BT)^3}\int_0^\infty R_{3\alpha}(E)E^2e^{-E/(k_BT)}dE \end{array}$ (26)

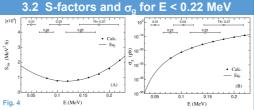
3.1 Analytic form of reaction rates



 $s_0 \left[\left(1 + \frac{5T_a}{36E_0} \right) + s_1 E_0 \left(1 + \frac{35T_a}{36E_0} \right) + s_2 E_0^2 \left(1 + \frac{89T_a}{36E_0} \right) \right], E_0 = (\pi \eta_0 T_a)^{2/3}. (30)$ Resonant contribution: $\langle R_{3a} \rangle_R = \frac{9\sqrt{3}\pi^3}{4} \frac{N_A^2 h^5}{m_N^3} \frac{\Gamma_{\gamma}(0_2^+)}{(k_B T)^3} \exp\left(-\frac{E(0_2^+)}{k_B T}\right)$ (31)

 $\approx 7.605 \times 10^{-9} \frac{\Gamma_{\gamma}(0_{2}^{+})}{\tau^{3}} \exp\left(-\frac{11.605 E(0_{2}^{+})}{\tau^{3}}\right)$ (32)

REACLIB [18] $\langle R_{3\alpha} \rangle = \sum_{i} \exp \left(a_{i0} + a_{i1}/T_9 + a_{i2}/T_9^{1/3} + a_{i3}T_9^{1/3} + a_{i4}T_9 + a_{i5}T_9^{5/3} + a_{i6}ln(T_9) \right)$ (33)



Compared with the cross sections in Fig.4(B), the energy dependence of the S-

The reaction rates below T₉= 0.02 are generated from the extrapolated S-factors However, I do not think that the derived rates have large uncertainties, because the S-factors do not seem to have the considerable variation to energies.

Table 1		Coefficients					
	s_0	s_1	s_2	η_0	а	$E(0_{2}^{+})$	$\Gamma_{\gamma}(0^+_2)$
	(MeV2b)	(MeV^{-1})	(MeV^{-2})	$(MeV^{1/2})$	(MeV^{-1})	(MeV)	(meV)
	2.570×10 ⁴	-12.12	51.46	4.153	5.0	0.3796	3.908

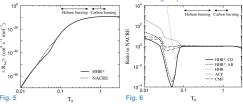
3.3 Comparison between reaction rates

The derived rates are consistent with NACRE for 0.08 < To

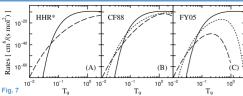
In contrast, the reaction rates are reduced by 10^4 at $T_9 \approx 0.05$. This is caused by the non-adiabatic description of $^8\mathrm{Be}$ break-up

 σ_n is reduced from that of ACF and CMF with the sequential process at E = $_{0.18}^{9}$ MeV (Fig. 3). Due to the strong influence of 0 $^{\circ}_{2}$, the difference in σ_{g} for E > 0.2 MeV cannot be found in the reaction rates.

Astrophysical impact of the direct triple- α process is expected to be small, because the difference is found before helium burning temperatures



Translation into REACLIB



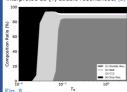
- The reaction rates are defined in Eq. (33). HHR* does not have a c
- The reaction rates are defined in Eq. (53), rain a cooling large in Fig.7). This means that the statistically generated 8B is broken-up immediately by the
- Third α-particle before its lifetime at T₉ ≈ 0.05.

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Table 2	i	a_{i0}	a_{i1}	a_{i2}	a_{i3}	a_{i4}	a_{i5}	a_{i6}	(%)	
	1	2.296	0.	-37.25	0.	-1.4	0.	-13/6	19%	
	2	-17.33	-4.405	0.	0.	0.	0.	-3.	n/a	

4. Three components in NACRE



The integrals of E₁ and E₂ in Eq. (34) can be divided into four terms. They are interpreted as (1) double resonances, (2) non-resonant sequential process, (3



, (3) 12C resonance, (4) non-resonant component. The composition ratio is shown in Fig. 8. The component of (3) appears to be small over the entire region. Therefore

the standard reaction rates for 0 below T₉ = 3 are found to generally consist of three components. The contribution of (2) is found to dominate the rates for $0.03 < T_9 < 0.07$

However, this term may not se

5. $\alpha + \alpha$ and 3α interaction

Two types of interaction potentials are used in Eq. (3). $V_{\alpha\alpha}^{N}(\bar{x}_{i}) = (V_{c0}\mathbf{P_{0}} + V_{c2}\mathbf{P_{2}})e^{-(\bar{x}_{i}/a)^{2}} + V_{0}e^{-(\bar{x}_{i}/b)^{2}},$ $V_{3\alpha}^{l}(\rho) = v_{3\alpha,l} \exp \left[-\left(\frac{\rho}{r_{3\alpha}}\right)^{2}\right]$ (35)

AB: shallow [20,9,13]

AB: shallow [20,9,15] CD: core-deep, folding potentials [21,22], determined by phase shifts & α -width AB & CD have the 3α potentials to reproduce the energies of 0^*_2 & 2^*_1 in 12 C.

6. Summary

In the present poster, I have described the direct triple- α process with non-adiabatic Faddeev HHR*, and I have discussed the reaction rates. The derived rates are expressed in a simple analytic form, and they are ed into REACLIB format.

Direct triple- α contribution is confirmed to be 10-15 10-3 pb order in the photo-disintegration cross sections of ${}^{4}\text{CC}(2^{\bullet}, \rightarrow 0^{\bullet})$ for 0.15 < E < 0.35 MeV. This is far below the values predicted by ACF and CMF that include the assumed long resonant tail of 0^{\bullet}_{2} . In spite of the large difference between the non-adiabatic and adiabatic cross sections, the derived rates are concordant with NACRE at the helium burning temperatures.

The difference in σ_g below E < 0.2 MeV can be seen in the rates for T_9 < 0.07. From the comparison among the calculations, I have found reaction rates could be reduced by about 10-4 at T_0 = 0.05, becau non-adiabatic description of Be break-up and continuum states.

The derived reaction rates have been expressed in the simple an rm as Eqs. (28)_(32) with Table 1. The present rates do not have the component corresponding to the non-resonant sequential process between α+⁸Be. I, therefore, find that this component could be at least eliminated by hand, to update the reaction rates in NACRE & REACLIB

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