Effects of entrance channel on fusion probability in hot fusion reactions *

Long Zhu(祝龙)^{1;1)} Jun Su(苏军)¹ Ching-Yuan Huang(黄庆元)¹ Feng-Shou Zhang(张丰收)^{2,3;2)}

¹ Sino-French Institute of Nuclear Engineering and Technology, Sun Yat-sen University, Zhuhai 519082, China

² The Key Laboratory of Beam Technology and Material Modification of Ministry of Education, College of Nuclear Science and

Technology, Beijing Normal University, Beijing 100875, China

 3 Beijing Radiation Center, Beijing 100875, China

Abstract: Within the framework of the dinuclear system (DNS) model, the fusion reactions leading to the compound nuclei 274 Hs^{*} and 286 Cn^{*} are investigated. The fusion probability as a function of DNS excitation energy is studied. The calculated results are in good agreement with the available experimental data. The obtained results show that the fusion probabilities are obviously enhanced for the reactions located at high place in potential energy surface, although these reactions may have small values of mass asymmetry. It is found that the enhancement is due to the large potential energy of the initial DNS.

Keywords: fusion probability, entrance channel effects, hot fusion reactions, dinuclear system model

PACS: 25.60.Pj, 25.70.Jj **DOI:** 10.1088/1674-1137/40/12/124105

1 Introduction

Production of superheavy nuclei (SHN) is one of most important research areas in nuclear physics. Heavy ion fusion reactions have been extensively investigated in synthesis of SHN experimentally and theoretically. The SHN with Z = 102 - 118 have been synthesized in the ²⁰⁸Pb- and ²⁰⁹Bi-based cold fusion reactions [1, 2] and ⁴⁸Ca-induced hot fusion reactions [3–5]. Recently, an experiment has been performed to synthesize the superheavy element (SHE) Z = 120 using the reaction ²⁴⁴Pu(⁵⁸Fe,xn)^{302-x}120 at FLNR (Dubna) [6]. Unfortunately, no correlated decay chains were observed. Cap et al. predicted that it is impossible to synthesize SHEs with Z = 122 and 124 in the fusion of symmetric systems [7]. Hence, the choice of more mass asymmetric reactions would be preferable. Choosing favorable reactions is very important for sythesizing SHEs with Z > 118. Therefore, it is worth investigating the mechanism of heavy ion fusion reactions theoretically.

The evaporation residual (ER) cross sections of fusion reactions strongly depend on the projectile-target combination. A lot of work has been done to investigate the entrance channel effects on production of SHN [8–15], including the reaction Q value and orientation effects on the ER cross sections [16–19]. However, the mechanism of entrance channel effects on productions of SHN, especially entrance channel effects of the fusion probability $(P_{\rm CN})$, is still not clear. Fusion probability is the least well known quantity that determines the ER cross sections. It is known that the fusion probability of massive nuclei strongly depends on the mass (charge) asymmetry $\eta = (A_1 - A_2)/(A_1 + A_2) \ (\eta_Z = (Z_1 - Z_2)/(Z_1 + Z_2)),$ where A_1, A_2 and Z_1, Z_2 are the mass and charge numbers of two fragments. Usually, $P_{\rm CN}$ is larger for a mass asymmetric reaction than for a symmetric one, because of the lower inner fusion barrier $B_{\rm fus}$ and the higher quasifission barrier B_{qf} [20]. But the opposite behavior can be seen for the reactions ${}^{124}Sn + {}^{92}Zr$ and ${}^{86}Kr + {}^{130}Xe$ in Ref. [9]. Therefore, it is desirable to get more detailed studies about the effects of entrance channel on $P_{\rm CN}$ in the synthesis of SHN.

Comparison of $P_{\rm CN}$ in different mass asymmetry reactions leading to the same compound nucleus (CN) allows us to analyze the entrance channel effects on the fusion-fission reaction mechanism in collisions of massive nuclei. The DNS model has been successfully used in investigating the synthesis mechanism of SHN [20–31]. In this work, we focus on the entrance channel effects of $P_{\rm CN}$ in synthesis of SHN within the framework of the DNS

Received 6 June 2016

^{*} Supported by Natural Science Foundation of Guangdong Province China (2016A030310208). National Natural Science Foundation of China (11605296, 11405278, 11505150, 11635003), Fundamental Research Funds for the Central Universities (15 lgpy 30) and China Postdoctoral Science Foundation (2015M582730)

¹⁾ E-mail: zhulong@mail.sysu.edu.cn

²⁾ E-mail: fszhang@bnu.edu.cn

^{©2016} Chinese Physical Society and the Institute of High Energy Physics of the Chinese Academy of Sciences and the Institute of Modern Physics of the Chinese Academy of Sciences and IOP Publishing Ltd

model. The ER cross sections and $P_{\rm CN}$ for the reactions ${}^{26}{\rm Mg} + {}^{248}{\rm Cm}$, ${}^{36}{\rm S} + {}^{238}{\rm U}$, and ${}^{48}{\rm Ca} + {}^{226}{\rm Ra}$ are investigated. The fusion probabilities leading to the compound nucleus ${}^{286}{\rm Cn}$ in various reactions ${}^{48}{\rm Ca} + {}^{238}{\rm U}$, ${}^{52}{\rm Ca} + {}^{234}{\rm U}$, ${}^{53}{\rm Ca} + {}^{233}{\rm U}$, and ${}^{55}{\rm Sc} + {}^{231}{\rm Pa}$ are investigated.

The article is organized as follows. In Section 2, we describe the DNS model. The results and discussion of $P_{\rm CN}$ are presented in Section 3. Finally, we summarize the main results in Section 4.

2 Description of the model

The ER cross sections of SHN can be written as a sum over all partial waves J,

$$\sigma_{\rm ER}(E_{\rm c.m.}) = \frac{\pi \hbar^2}{2\mu E_{\rm c.m.}} \sum_J (2J+1)T(E_{\rm c.m.},J) \\ \times P_{\rm CN}(E_{\rm c.m.},J)W_{\rm sur}(E_{\rm c.m.},J).$$
(1)

Here, T, $P_{\rm CN}$, and $W_{\rm sur}$ are transmission probability, fusion probability, and survival probability, respectively. $E_{\rm c.m.}$ and J are the incident energy and incident angular momentum in the center of mass frame, respectively. In this work, the transmission probability T in the capture process is calculated by empirical coupled-channel model [29], in which the Woods-Saxon potential is used for nuclear potential, and its parameters $V_0 = 80$ MeV and a = 0.68 fm, are adopted.

The fusion probability is investigated within DNS model. In this work, the diffusion process is treated along proton and neutron degrees of freedom. The probability distribution function $P(Z_1, N_1, E_1, t)$ for fragments 1 with proton number Z_1 , neutron number N_1 and excitation energy E_1 at time t can be obtained by solving the following master equations [22],

$$\begin{split} & \frac{\mathrm{d}P(Z_1,N_1,E_1,t)}{\mathrm{d}t} \\ &= \sum_{Z_1'} W_{Z_1,N_1;Z_1',N_1}(t) [d_{Z_1,N_1} P(Z_1',N_1,E_1',t) \\ &- d_{Z_1',N_1} P(Z_1,N_1,E_1,t)] \\ &+ \sum_{N_1'} W_{Z_1,N_1;Z_1,N_1'}(t) [d_{Z_1,N_1} P(Z_1,N_1',E_1',t) \\ &- d_{Z_1,N_1'} P(Z_1,N_1,E_1,t)] \\ &- A_{\mathrm{qf}}(\Theta(t)) P(Z_1,N_1,E_1,t). \end{split}$$

Here $W_{Z_1,N_1;Z'_1,N_1}$ $(W_{Z_1,N_1;Z_1,N'_1})$ denotes the mean transition probability from the channel (Z_1, N_1, E_1) to (Z'_1, N_1, E'_1) [or (Z_1, N_1, E_1) to (Z_1, N'_1, E'_1)], and d_{Z_1,N_1} is the microscopic dimension corresponding to the macroscopic state (Z_1, N_1, E_1) . In the DNS model, we consider the process of only one nucleon transfer. The sum is taken over all possible proton and neutron numbers that fragment 1 may take. $\Lambda_{\rm qf}$ is the quasifission rate, which describes the evolution of DNS system along relative distance R. The detailed description of $\Lambda_{\rm qf}$ can be seen in Ref. [17].

The local excitation energy E_1 in Eq. (2) is related to the excitation energy of the composite system at the initial state E_{DNS}^* and the driving potentials of the injection point of the DNS. E_1 is given by

$$E_{1} = E_{\text{DNS}}^{*} - U(Z_{1}, N_{1}, Z_{2}, N_{2}, J) + U(Z_{\text{p}}, N_{\text{p}}, Z_{\text{t}}, N_{\text{t}}, J) - \frac{(J - M)^{2}}{2\zeta_{\text{rel}}} - \frac{M^{2}}{2\zeta_{\text{int}}}, \quad (3)$$

where $U(Z_1, N_1, Z_2, N_2)$ and $U(Z_p, N_p, Z_t, N_t)$ are the potential energies of the configuration $(Z_1, N_1; Z_2, N_2)$ and the DNS at the injection point. M denotes the intrinsic angular momentum derived from the dissipation of the relative angular momentum, and ζ_{int} is the corresponding moment of inertia. J denotes the initial angular momentum. $\zeta_{\rm rel}$ is the relative moment of inertia of the DNS, which is given by $\zeta_{\rm rel} = \mu R_{\rm m}^2$. The local excitation energy means the excitation energy of the DNS for a specific configuration $(Z_1, N_1; Z_2, N_2)$. E^*_{DNS} is converted from the relative kinetic energy loss, which is related to the incident energy and the minimum of the well bottom of the nucleus-nucleus potential in the entrance channel $V(R_{\rm m})$ [32]. Here, $R_{\rm m}$ is the distance of two colliding nuclei located at the bottom of the potential pocket. E^*_{DNS} can be written as,

$$E_{\rm DNS}^* = E_{\rm c.m.} - V(R_{\rm m}) - \frac{(J\hbar)^2}{2\zeta_{\rm rel}}.$$
 (4)

The potential energy surface (PES) of the DNS is given by

$$U(Z_1, N_1, R_m, J) = B(Z_1, N_1) + B(Z_2, N_2) - B(Z, N) - V_{\rm rot}^{\rm CN}(J) + V_{\rm CN}(Z_1, N_1, R_m, J).$$
(5)

 $B(Z_i, N_i)$ (i = 1, 2) and B(Z, N) are the ground state binding energies of the fragment *i* and compound nucleus $(Z = Z_1 + Z_2; N = N_1 + N_2)$, respectively. $V_{\rm CN}$ is the interaction potential of two fragments. The details of $V_{\rm CN}$ are given in Ref. [21]. The $Q_{\rm gg}$ value of forming a configuration $(Z_1, N_1; Z_2, N_2)$ is defined as

$$Q_{\rm gg} = B_{\rm DNS}(Z_{\rm p}, N_{\rm p}; Z_{\rm t}, N_{\rm t}) - B_{\rm DNS}(Z_1, N_1; Z_2, N_2).$$
(6)

Here, B_{DNS} is the binding energy of DNS system, which can be written as $B_{\text{DNS}}(Z_1, N_1; Z_2, N_2) = B(Z_1, N_1) + B(Z_2, N_2)$.

The fusion probability is expressed as follows:

$$P_{\rm CN}(E_{\rm c.m.},J) = \sum_{A_1=1}^{A_{\rm BG}} P(A_1, E_{\rm c.m.},J),$$
(7)

where $A_{\rm BG}$ (= $N_{\rm BG} + Z_{\rm BG}$) is the mass number of light fragment at Businaro-Gallone (B.G.) point.

The survival probability of emitting x neutrons can be written as

$$W_{\rm sur}(E_{\rm CN}^{*}, x, J) = P(E_{\rm CN}^{*}, x, J) \\ \times \prod_{i}^{x} [\frac{\Gamma_{\rm n}(E_{i}^{*}, J)}{\Gamma_{\rm n}(E_{i}^{*}, J) + \Gamma_{\rm f}(E_{i}^{*}, J)}].$$
(8)

Here E_{CN}^* , J are the excitation energy and the spin of the compound nucleus, respectively. $E_i^* (= E_{i-1}^* - B_{i-1}^n - 2T_{i-1})$ is the excitation energy before evaporation of the *i*th neutron. The detailed description of the width of the *i*th neutron emission and fission can be found in Ref. [29]. The realization probability $P(E_{\text{CN}}^*, x, J)$ is calculated as in Ref. [11]. In this work, the fission barrier before evaporating the *i*th neutron is obtained by

$$B_i^{\rm f}(E_i^*) = B_{\rm M}^{\rm f}(E_i^* = 0) \exp(-E_i^*/E_{\rm d}).$$
(9)

The microscopic shell correction energy $B_{\rm M}^{\rm f}(E_i^*=0)$ is taken from Ref. [33]. The damping factor $E_{\rm d}=20~{\rm MeV}$ is taken.

3 Results and discussion

3.1 Reactions to the compound nucleus ²⁷⁴Hs*

The DNS model is used to investigate the fusion reactions ${}^{26}Mg + {}^{248}Cm$, ${}^{36}S + {}^{238}U$, and ${}^{48}Ca + {}^{226}Ra$ leading to the same compound nucleus ${}^{274}Hs^*$.

Figure 1(a) shows the PES as functions of proton and neutron number of the fragment 1 for the DNS configurations in formation of the compound nucleus ²⁷⁴Hs^{*}. The red line indicates the minimum trajectory in the PES. We show injection points in the PES for the reactions ²⁶Mg + ²⁴⁸Cm, ³⁶S + ²³⁸U, and ⁴⁸Ca + ²²⁶Ra, which are all close to the red line. The minimum trajectory in the PES (U^{\min}) can be presented as a function of the mass asymmetry η , as shown in Fig. 1(b). The dashed arrow denotes the position of the B.G. point. The vertical solid arrow indicate the inner fusion barrier B_{fus} for the reaction ⁴⁸Ca + ²²⁶Ra. Usually, the inner fusion barrier B_{fus} and quasifission barrier B_{qf} play a main role in the competition between quasifission and complete fusion [9, 20].

Figure 2 shows the ER cross sections for the reactions ${}^{26}Mg + {}^{248}Cm$, ${}^{36}S + {}^{238}U$, and ${}^{48}Ca + {}^{226}Ra$. The calculated ER cross sections are in agreement with the experimental data within a factor of 1–3. For the reaction ${}^{36}S + {}^{238}U$, we predict that the maximal ER cross section of 4n channel is 11.6 pb, and the corresponding excitation energy equals $E^* = 41$ MeV.

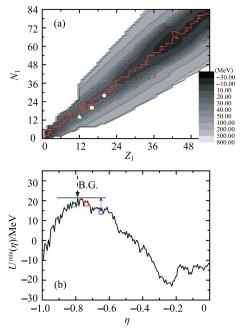


Fig. 1. (color online) (a) The PES as functions of Z_1 and N_1 of the fragment 1 for the DNS configurations leading to the compound nucleus 274 Hs^{*}. The red line indicates the minimum trajectory in the PES. Triangle, square, and circle represent the injection points for the reactions 26 Mg + 248 Cm, 36 S + 238 U, and 48 Ca + 226 Ra, respectively. (b) The driving potential, $U^{\min}(\eta)$, which is the line connecting the minimum trajectory as a function of mass asymmetry η . The dashed arrow denotes the position of the B.G. point. The vertical solid arrow indicates the inner fusion barrier for the reaction 48 Ca + 226 Ra.

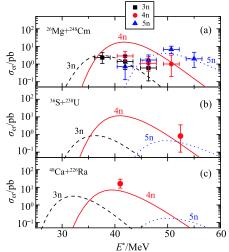


Fig. 2. (color online) Comparison of the calculated ER cross sections with the available experimental data [4, 34, 35] for the reactions ²⁶Mg + ²⁴⁸Cm (a), ³⁶S + ²³⁸U (b), and ⁴⁸Ca + ²²⁶Ra (c). The experimental data of 3n, 4n, and 5n channels are denoted by squares, circles, and triangles, respectively.

Figure 3 presents the dependence of the fusion probabilities on excitation energy of DNS for the reactions ${}^{26}Mg + {}^{248}Cm$, ${}^{36}S + {}^{238}U$, and ${}^{48}Ca + {}^{226}Ra$ in formation of same compound nucleus ${}^{274}Hs^*$. The $P_{\rm CN}$ is calculated with J = 0 [13]. The available experimental data for the reactions ${}^{26}Mg + {}^{248}Cm$ and ${}^{36}S + {}^{238}U$ are also shown. It can be seen from the experimental data that the fusion probabilities for the reaction ${}^{26}Mg + {}^{248}Cm$ are larger than those for the reaction ${}^{36}S + {}^{238}U$. The calculated results show the same behavior. One can see that the fusion probability decreases with the decreasing mass asymmetry for the reactions ${}^{26}Mg + {}^{248}Cm$, ${}^{36}S + {}^{238}U$, and ${}^{48}Ca + {}^{226}Ra$. The same trend can be seen in Ref. [11, 16]. Table 1 shows the quasifission barriers B_{qf} , inner fusion barriers B_{fus} , potential energy at the bottom of pocket $V(R_m)$, and Coulomb barriers V_B for different reactions. For the reactions ${}^{26}Mg + {}^{248}Cm$, ${}^{36}S + {}^{238}U$, and ${}^{48}Ca + {}^{226}Ra$, it is noticed that the B_{fus} increases and B_{qf} decreases with the decrease of mass asymmetry. This is the main reason for the behavior of P_{CN} observed above.

Table 1. Physical quantities (the quasifission barrier B_{qf} , the inner fusion barrier B_{fus} , $V(R_m)$, and Coulomb barrier V_B) for the indicated reactions. Angular momentum J = 0.

reactions	compound nuclei	$B_{\rm qf}/{ m MeV}$	$B_{\rm fus}/{ m MeV}$	$V(R_{ m m})/{ m MeV}$	$V_{\rm B}/{ m MeV}$
$^{26}Mg + ^{248}Cm$	274 Hs*	10.72		111.80	122.52
$^{36}S + ^{238}U$	274 Hs*	8.96	2.71	144.12	153.08
${}^{48}\text{Ca} + {}^{226}\text{Ra}$	274 Hs*	6.65	6.82	169.80	176.45
${}^{48}\text{Ca} + {}^{238}\text{U}$	$^{286}\mathrm{Cn}^*$	6.25	10.29	177.13	183.38
${}^{52}Ca + {}^{234}U$	$^{286}\mathrm{Cn}^*$	6.60	9.08	175.24	181.84
$^{53}Ca + ^{233}U$	$^{286}\mathrm{Cn}^{*}$	6.70	7.86	174.83	181.53
$^{55}Sc + ^{231}Pa$	$^{286}\mathrm{Cn}^{*}$	5.86	10.47	182.39	188.25

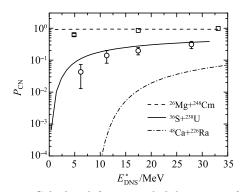


Fig. 3. Calculated fusion probabilities as a function of E_{DNS}^* (J = 0) for the reactions ${}^{26}\text{Mg} + {}^{248}\text{Cm}, {}^{36}\text{S} + {}^{238}\text{U}, \text{ and } {}^{48}\text{Ca} + {}^{226}\text{Ra}.$ Experimental values of P_{CN} obtained in Ref. [36] for the reactions ${}^{26}\text{Mg} + {}^{248}\text{Cm}$ and ${}^{36}\text{S} + {}^{238}\text{U}$ are denoted by squares and circles, respectively.

3.2 Fusion probabilities in formation of the compound nucleus ²⁸⁶Cn*

To clarify the entrance channel effects on $P_{\rm CN}$, we also apply the model to calculate the fusion probabilities in the reactions ${}^{48}\text{Ca} + {}^{238}\text{U}$, ${}^{52}\text{Ca} + {}^{234}\text{U}$, and ${}^{53}\text{Ca} + {}^{233}\text{U}$, and ${}^{55}\text{Sc} + {}^{231}\text{Pa}$. These reactions are all in formation of the same compound nucleus ${}^{286}\text{Cn}^*$.

Figure 4(a) shows the PES as functions of proton and neutron number of fragments 1 for DNS configurations leading to the compound nucleus ${}^{286}Cn^*$. The injection points for the reactions ${}^{53}Ca + {}^{233}U$, ${}^{52}Ca + {}^{234}U$, ${}^{48}Ca$

+ ²³⁸U, and ⁵⁵Sc + ²³¹Pa are denoted by triangle, circle, square, and star, respectively. It can be seen that the injection points for the reactions ⁵³Ca + ²³³U and ⁵⁵Sc + ²³¹Pa are located at quite a high place. The reason is that the nuclei ⁵³Ca and ⁵⁵Sc are radioactive. The minimum trajectory in the PES (U^{\min}) can be presented as a function of the mass asymmetry η , as shown in Fig. 4(b).

Figure 5(a) shows the comparison of capture cross section between results in this work and the experimental data [37–39] for the ⁴⁸Ca + ²³⁸U fusion reaction. The calculated result is in good agreement with the experimental data from Ref. [37] over the whole incident energy region. It can be seen that calculated result also shows good agreement with all experimental data measured by Nishio et al [39] around and above the Coulomb barrier. Figure. 5(b) shows a comparison of ER cross sections between experimental data and calculated results. Within the error bars, the experimental data [5] are reproduced rather well, especially for the 4n channel.

Figure 6(a) shows the fusion probabilities for the reactions ${}^{48}\text{Ca} + {}^{238}\text{U}$, ${}^{52}\text{Ca} + {}^{234}\text{U}$, and ${}^{53}\text{Ca} + {}^{233}\text{U}$. For these reactions, the difference of mass asymmetry is induced by neutron number. It can be seen that P_{CN} for the reaction ${}^{53}\text{Ca} + {}^{233}\text{U}$ is much larger than that for the reaction ${}^{48}\text{Ca} + {}^{238}\text{U}$ at large E_{DNS}^* range, although ${}^{53}\text{Ca} + {}^{233}\text{U}$ shows smaller mass asymmetry. At high E_{DNS}^* region, the fusion probability for the reaction ${}^{52}\text{Ca} + {}^{234}\text{U}$ is lower than that for the reaction ${}^{48}\text{Ca} + {}^{238}\text{U}$,

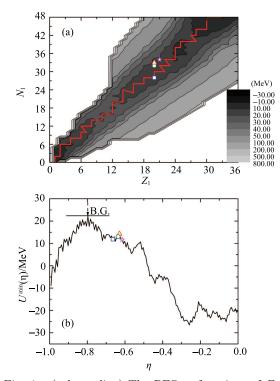


Fig. 4. (color online) The PES as functions of Z_1 and N_1 of fragments 1 for the DNS configurations leading to the compound nucleus ²⁸⁶Cn^{*}. The injection points for the reactions ⁵³Ca + ²³³U, ⁵²Ca + ²³⁴U, ⁴⁸Ca + ²³⁸U, and ⁵⁵Sc + ²³¹Pa are denoted by triangle, circle, square, and star, respectively.

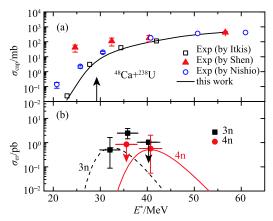


Fig. 5. (color online) (a) Capture cross sections for the reaction ${}^{48}\text{Ca} + {}^{238}\text{U}$. The experimental data are denoted by squares [37], triangles [38], and circles [39]. The solid line denotes the calculated result in this work. The vertical arrow shows the position of Coulomb barrier. (b) ER cross sections for the reaction ${}^{48}\text{Ca} + {}^{238}\text{U}$. The dashed line and solid line denote the calculated ER cross sections of the 3n and 4n channels, respectively. The experimental data [5] of the 3n and 4n channels are denoted by solid squares and circles, respectively.

while the opposite behavior is shown at low E_{DNS}^* . In Fig. 6(b), we notice that the fusion probability of the reaction ${}^{55}\text{Sc} + {}^{231}\text{Pa}$ is larger than that of the reaction ${}^{48}\text{Ca} + {}^{238}\text{U}$, although the reaction ${}^{55}\text{Sc} + {}^{231}\text{Pa}$ has a smaller mass asymmetry (induced by proton number). It is already known that the barriers B_{fus} and B_{qf} play crucial roles in complete fusion [9]. Usually, in formation of the same compound nucleus, the fusion probability is larger for the reaction with smaller B_{fus} and larger B_{qf} , which means a larger fusion probability for the reaction ${}^{48}\text{Ca} + {}^{238}\text{U}$ than that for the reaction ${}^{55}\text{Sc} + {}^{231}\text{Pa}$ (see Table 1). However, the opposite behavior is shown. Therefore, some other physical quantities need to be taken into account, especially for radioactive beam induced hot fusion reactions.

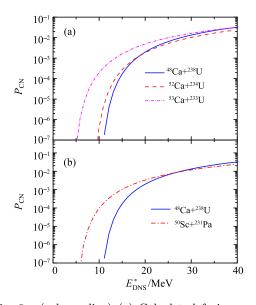


Fig. 6. (color online) (a) Calculated fusion probabilities as a function of E_{DNS}^* (J = 0) for the reactions ${}^{48}\text{Ca} + {}^{238}\text{U}$, ${}^{52}\text{Ca} + {}^{234}\text{U}$, and ${}^{53}\text{Ca} + {}^{233}\text{U}$ with different mass asymmetry induced by neutron number. (b) Calculated fusion probabilities as a function of E_{DNS}^* (J = 0) for the reactions ${}^{48}\text{Ca} + {}^{238}\text{U}$ and ${}^{55}\text{Sc} + {}^{231}\text{Pa}$ with different mass asymmetry induced by proton number.

As we discussed in Fig. 4, the injection point in the PES for the reaction ${}^{52}\text{Ca} + {}^{234}\text{U}$, ${}^{53}\text{Ca} + {}^{233}\text{U}$, and ${}^{55}\text{Sc} + {}^{231}\text{Pa}$ is located at a higher place than the reaction ${}^{48}\text{Ca} + {}^{238}\text{U}$. To clarify the influence of potential energy of initial DNS on the fusion probability, we define the difference, $\Delta U = U^{\text{inj}}(Z_1, N_1) - U^{\text{min}}(\eta)$, between the potential energy at the injection point $U^{\text{inj}}(Z_1, N_1)$ (as shown in Fig. 4(a)) and the driving potential $U^{\text{min}}(\eta)$ with corresponding value of mass asymmetry at the minimum trajectory (as shown in Fig. 4(b)). The difference, $\Delta V(R_{\text{m}}) = V^{\text{inj}}(R_{\text{m}}) - V^{\text{min}}(R_{\text{m}})$, is also defined. Here,

the superscripts "inj" and "min" denote the initial combinations at the injection points and configurations with the same mass asymmetry at the minimum trajectory, respectively. The difference of $Q_{\rm gg}$ value, in formation of DNS configuration $(Z_1, N_1; Z_2, N_2)$, between initial DNS configurations at the injection points and corresponding configurations with the same mass asymmetry at minimum trajectory can be written as $\Delta Q_{\rm gg} = B_{\rm DNS}^{\rm inj} - B_{\rm DNS}^{\rm min}$. Therefore, $\Delta U = \Delta Q_{\rm gg} + \Delta V(R_{\rm m})$.

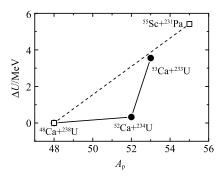


Fig. 7. ΔU for the combinations leading to the same compound nucleus ²⁸⁶Cn^{*}. The circles and squares denote the reactions with different mass asymmetry induced by neutron number and proton number, respectively. The lines are used to guide the eye.

We show ΔU for the reactions ⁴⁸Ca + ²³⁸U, ⁵²Ca + ²³⁴U, ⁵³Ca + ²³³U, and ⁵⁵Sc + ²³¹Pa in Fig. 7. It can be seen that ΔU for the reactions ⁵³Ca + ²³³U and ⁵⁵Sc + ²³¹Pa is much larger than that for the reaction ⁴⁸Ca + ²³⁸U. Apart from the local excitation energy of forming a DNS configuration (Z_1 , N_1 ; Z_2 , N_2) in the corresponding configuration with same mass asymmetry located at

References

- S. Hofmann and G. Münzenberg, Rev. Mod. Phys. **72**: 733 (2000);
 S. Hofmann, F. P. Heßberger, D. Ackermann, G. Münzenberg, S. Antalic, P. Cagarda, B. Kindler, J. Kojouharova, M. Leino, B. Lommel, R. Mann, A. G. Popeko, S. Reshitko, S. Saro, J. Uusitalo, and A. V. Yeremin, Eur. Phys. J. A, **14**: 147–157 (2002)
- 2 K. Morita, K. Morimoto, D. Kaji, H. Haba, K. Ozeki, Y. Kudou, T. Sumita, Y. Wakabayashi, A. Yoneda, K. Tanaka, S. Yamaki, R. Sakai, T. Akiyama, S. Goto, H. Hasebe, M. Huang, T. Huang, E. Ideguchi, Y. Kasamatsu, K. Katori, Y. Kariya, H. Kikunaga, H. Koura, H. Kudo, A. Mashiko, K. Mayama, S. Mitsuoka, T. Moriya, M. Murakami, H. Murayama, S. Namai, A. Ozawa, N. Sato, K. Sueki, M. Takeyama, F. Tokanai, T. Yamaguchi, and A. Yoshida, J. Phys. Soc. Jpn., **81**: 103201 (2012)
- 3 Yu. Ts. Oganessian, J. Phys. G: Nucl. Part. Phys., 34: R165-R242 (2007)
- 4 Yu. Ts. Oganessian, V. K. Utyonkov, F. Sh. Abdullin, S. N. Dmitriev, R. Graeger, R. A. Henderson, M. G. Itkis, Yu. V. Lobanov, A. N. Mezentsev, K. J. Moody, S. L. Nelson, A. N. Polyakov, M. A. Ryabinin, R. N. Sagaidak, D. A. Shaughnessy,

the minimum trajectory, some extra excitation energy $(= \Delta U)$ will be released in the reactions ${}^{52}\text{Ca} + {}^{234}\text{U}$, ${}^{53}\text{Ca} + {}^{233}\text{U}$, and ${}^{55}\text{Sc} + {}^{231}\text{Pa}$, as shown in Eq. (3). The extra excitation energies in the reactions ${}^{53}\text{Ca} + {}^{233}\text{U}$ and ${}^{55}\text{Sc} + {}^{231}\text{Pa}$ are much larger than those in the other reactions. Therefore, with the same E_{DNS}^* , in formation of the same DNS configuration $(Z_1, N_1; Z_2, N_2)$ the local excitation energies E_1 for the reactions ${}^{53}\text{Ca} + {}^{233}\text{U}$ and ${}^{55}\text{Sc} + {}^{231}\text{Pa}$ are larger than others. As we known, the fusion probability is sensitive to the value of the excitation energy. The P_{CN} usually increases with the increasing E_1 [21]. Therefore, the enhancement of fusion probability for the reactions ${}^{53}\text{Ca} + {}^{231}\text{Pa}$ are mainly due to large ΔU .

4 Conclusions

In this article, we have investigated the entrance channel effects of fusion probability within the DNS model. The reactions ${}^{26}Mg + {}^{248}Cm$, ${}^{36}S + {}^{238}U$, ${}^{48}Ca$ + ²²⁶Ra, ⁵³Ca + ²³³U, ⁵⁵Sc + ²³¹Pa, ⁴⁸Ca + ²³⁸U, and ${}^{52}Ca + {}^{234}U$ leading to the compound nuclei ${}^{274}Hs^*$ and $^{286}\mathrm{Cn}^*$ have been studied. It has been claimed that the fusion probability is enhanced for a mass asymmetric reaction [20]. However, our calculations show that the fusion probabilities as a function of excitation energy of DNS for the reactions ${}^{53}\text{Ca} + {}^{233}\text{U}$ and ${}^{55}\text{Sc} + {}^{231}\text{Pa}$ are larger than that for the ${}^{48}Ca + {}^{238}U$ reaction, although the mass asymmetry of ${}^{48}Ca + {}^{238}U$ is larger. The enhancement of fusion probability is mainly due to large value of potential energies of initial DNS. For future work, the capture cross sections and survival probabilities also need to be considered in finding favorable radioactive induced hot fusion reactions to synthesize SHN.

I. V. Shirokovsky, M. A. Stoyer, N. J. Stoyer, V. G. Subbotin, K. Subotic, A. M. Sukhov, Yu. S. Tsyganov, A. Türler, A. A. Voinov, G. K. Vostokin, P. A. Wilk, and A. Yakushev, Phys. Rev. C, **87**: 034605 (2013)

- 5 Yu. Ts. Oganessian, V. K. Utyonkov, Yu. V. Lobanov, F. Sh. Abdullin, A. N. Polyakov, I. V. Shirokovsky, Yu. S. Tsyganov, G. G. Gulbekian, S. L. Bogomolov, B. N. Gikal, A. N. Mezentsev, S. Iliev, V. G. Subbotin, A. M. Sukhov, A. A. Voinov, G. V. Buklanov, K. Subotic, V. I. Zagrebaev, M. G. Itkis, J. B. Patin, K. J. Moody, J. F. Wild, M. A. Stoyer, N. J. Stoyer, D. A. Shaughnessy, J. M. Kenneally, P. A. Wilk, R. W. Lougheed, R. I. Il'kaev, and S. P. Vesnovskii, Phys. Rev. C, **70**: 064609 (2004)
- 6 Yu. Ts. Oganessian, V. K. Utyonkov, Yu. V. Lobanov, F. Sh. Abdullin, A. N. Polyakov, R. N. Sagaidak, I. V. Shirokovsky, Yu. S. Tsyganov, A. A. Voinov, A. N. Mezentsev, V. G. Subbotin, A. M. Sukhov, K. Subotic, V. I. Zagrebaev, S. N. Dmitriev, R. A. Henderson, K. J. Moody, J. M. Kenneally, J. H. Landrum, D. A. Shaughnessy, M. A. Stoyer, N. J. Stoyer, and P. A. Wilk, Phys. Rev. C, **79**: 024603 (2009)
- 7 T. Cap, K. Siwek-Wilczyńska, J. Wilczyński, Phys. Lett. B,

 $\textbf{736:} \ 478\text{-}481 \ (2014)$

- 8 A. K. Nasirov, G. Giardina, G. Mandaglio, M. Manganaro, F. Hanappe, S. Heinz, S. Hofmann, A. I. Muminov, and W. Scheid, Phys. Rev. C, **79**: 024606 (2009)
- 9 G. Fazio, G. Giardina, G. Mandaglio, R. Ruggeri, A. I. Muminov, A. K. Nasirov, Yu. Ts. Oganessian, A. G. Popeko, R. N. Sagaidak, A. V. Yeremin, S. Hofmann, F. Hanappe, and C. Stodel, Phys. Rev. C, **72**: 064614 (2005)
- 10 K. Kim, Y. Kim, A. K. Nasirov, G. Mandaglio, and G. Giardina, Phys. Rev. C, **91**: 064608 (2015)
- 11 J. Hong, G. G. Adamian, and N. V. Antonenko, Phys. Rev. C, 92: 014617 (2015)
- 12 A. K. Nasirov, G. Mandaglio, G. Giardina, A. Sobiczewski, and A. I. Muminov, Phys. Rev. C, 84: 044612 (2011)
- 13 V. Zagrebaev and W. Greiner, Phys. Rev. C, 78: 034610 (2008)
- 14 C. W. Shen, Y. Abe, D. Boilley, G. Kosenko, and E. G. Zhao, Int. J. Mod. Phys. E, **17**: 66 (2008)
- 15 N. Wang, J. Tian, and W. Scheid, Phys. Rev. C, 84: 061601(R) (2011)
- 16 Z. H. Liu and J. D. Bao, Phys. Rev. C, 74: 057602 (2006)
- 17 L. Zhu, Z. Q. Feng, C. Li, and F. S. Zhang, Phys. Rev. C, 90: 014612 (2014)
- 18 A. S. Umar, V. E. Oberacker, J. A. Maruhn, and P. -G. Reinhard, Phys. Rev. C, 81: 064607 (2010)
- 19 K. Nishio, H. Ikezoe, S. Mitsuoka, I. Nishinaka, Y. Nagame, Y. Watanabe, T. Ohtsuki, K. Hirose, and S. Hofmann, Phys. Rev. C, 77: 064607 (2008)
- 20 G. G. Adamian, N. V. Antonenko, W. Scheid, and V. V. Volkov, Nucl. Phys. A, 633: 409 (1998)
- 21 Z. Q. Feng, G. M. Jin, J. Q. Li, and W. Scheid, Phys. Rev. C, 76: 044606 (2007)
- 22 Z. Q. Feng, G. M. Jin, and J. Q. Li, Nucl. Phys. A, 836: 82 (2010)
- 23 M. H. Huang, Z. G. Gan, X. H. Zhou, J. Q. Li, and W. Scheid, Phys. Rev. C, 82: 044614 (2010)
- 24 G. G. Adamian, N. V. Antonenko, and W. Scheid, Phys. Rev. C, 68: 034601 (2003)
- 25 G. G. Adamian, N. V. Antonenko, and W. Scheid, Phys. Rev. C, 69: 011601(R) (2004)
- 26 X. J. Bao, Y. Gao, J. Q. Li and H. F. Zhang, Phys. Rev. C, 92: 034612 (2015)
- 27 X. J. Bao, Y. Gao, J. Q. Li and H. F. Zhang, Phys. Rev. C,

93: 044615 (2016)

- 28 N. Wang, E. G. Zhao, W. Scheid, and S. G. Zhou, Phys. Rev. C, 85: 041601(R) (2012)
- 29 Z. Q. Feng, G. M. Jin, F. Fu, and J. Q. Li, Nucl. Phys. A, 771: 50 (2006)
- 30 G. G. Adamian, N. V. Antonenko, and W. Scheid, Phys. Rev. C 69: 014607 (2004); 69: 044601 (2004)
- 31 G. G. Adamian, N. V. Antonenko, and W. Scheid, Nucl. Phys. A, 678: 24 (2000)
- 32 G. Wolschin and W. Nörenberg, Z. Phys. A, **284**: 209 (1978)
- 33 P. Möller, J. R. Nix, W. D. Myers, and W. J. Swiatecki, At. Data Nucl. Data Tables, 59: 185 (1995)
- 34 J. Dvorak, W. Brüchle, M. Chelnokov, Ch.E. Düllmann, Z. Dvorakova, K. Eberhardt, E. Jäger, R. Krücken, A. Kuznetsov, Y. Nagame, F. Nebel, K. Nishio, R. Perego, Z. Qin, M. Schädel, B. Schausten, E. Schimpf, R. Schuber, A. Semchenkov, P. Thörle, A. Türler, M. Wegrzecki, B. Wierczinski, A. Yakushev, and A. Yeremin, Phys. Rev. Lett., **100**: 132503 (2008)
- 35 R. Graeger, D. Ackermann, M. Chelnokov, V. Chepigin, Ch. E. Düllmann, J. Dvorak, J. Even, A. Gorshkov, F. P. Heßberger, D. Hild, A. Hübner, E. Jäger, J. Khuyagbaatar, B. Kindler, J. V. Kratz, J. Krier, A. Kuznetsov, B. Lommel, K. Nishio, H. Nitsche, J. P. Omtvedt, O. Petrushkin, D. Rudolph, J. Runke, F. Samadani, M. Schädel, B. Schausten, A. Türler, A. Yakushev, and Q. Zhi, Phys. Rev. C, **81**: 061601(R) (2010)
- 36 I. M. Itkis, E. M. Kozulin, M. G. Itkis, G. N. Knyazheva, A. A. Bogachev, E. V. Chernysheva, L. Krupa, Yu. Ts. Oganessian, V. I. Zagrebaev, A. Ya. Rusanov, F. Goennenwein, O. Dorvaux, L. Stuttgé, F. Hanappe, E. Vardaci, E. de Goés Brennand, Phys. Rev. C, 83: 064613 (2011)
- 37 M. G. Itkis, A. A. Bogatchev, I. M. Itkis, M. Jandel, J. Kliman, G. N. Kniajeva, N. A. Kondratiev, I. V. Korzyukov, E. M. Kozulin, L. Krupa, Yu. Ts. Oganessian, I. V. Pokrovski, V. A. Ponomarenko, E. V. Prokhorova, A. Ya. Rusanov, V. M. Voskresenski, A. A. Goverdovski, F. Hanappe, T. Materna, N. Rowley, L. Stuttge, G. Giardina, and K. J. Moody, J. Nucl. Radiochem. Sci., **3**: 57 (2002)
- 38 W. Q. Shen et al., Phys. Rev. C, 36: 115 (1987)
- 39 K. Nishio, S. Mitsuoka, I. Nishinaka, H. Makii, Y. Wakabayashi, H. Ikezoe, K. Hirose, T. Ohtsuki, Y. Aritomo, and S. Hofmann, Phys. Rev. C, 86: 034608 (2012)