

Multidimensional fission model with a complex absorbing potentialGuillaume Scamps^{1,*} and Kouichi Hagino^{1,2,†}¹*Department of Physics, Tohoku University, Sendai 980-8578, Japan*²*Research Center for Electron Photon Science, Tohoku University, 1-2-1 Mikamine, Sendai 982-0826, Japan*

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We study the dynamics of multidimensional quantum tunneling by introducing a complex absorbing potential into a two-dimensional model for spontaneous fission. We first diagonalize the Hamiltonian with the complex potential to determine a resonance state as well as its lifetime. We then solve the time-dependent Schrödinger equation with such basis in order to investigate the tunneling path. We compare this method with the semiclassical method for multidimensional tunneling with imaginary time. A good agreement is found both for the lifetime and for the tunneling path.

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I. INTRODUCTION

Spontaneous fission is a typical example of multidimensional quantum tunneling, and its description has remained a challenge in nuclear theory. The first step for any calculation is to construct a potential energy surface in a multidimensional space. A standard approach for this is the macroscopic-microscopic method, with which the potential surface is constructed using the shell correction method of Strutinsky [1–3]. In recent years, a microscopic description based on a self-consistent mean-field theory has also been attempted [4–17]. Even though a microscopic understanding of fission phenomena is important, there have still been many open problems to be solved. For instance, the choice of relevant degrees of freedom is still under discussion [9] and a large uncertainty may arise from a choice of energy functional [18]. Moreover, a difficulty in constrained mean-field calculations has also been pointed out [3,19].

In order to calculate the fission lifetime, most of the calculations, with both the macroscopic and microscopic approaches, rely on the semiclassical approximation. That is, one often searches the least action path in a multidimensional space [1,7,9] or equivalently solves the Newtonian equations with the inverted potential [20–22].

In this paper, we investigate this problem from a different perspective. That is, we solve the time-dependent Schrödinger equation (TDSE) and monitor the time evolution of the wave function in a fully quantum mechanical manner. This method provides a good intuitive description for particle decays, and has been applied to systems where the decay width of a resonance state is about the same order of magnitude as the resonance energy [23–27].

In the previous applications of this method, because the TDSE has been solved by numerical integration, the time step had to be small with respect to the typical time scale of the process. The evolution was then restricted to the small time regime. In this paper, we propose a new method introducing a complex absorbing potential in the exterior region. The

complex absorbing potential has been used in time-dependent calculations [28–32] in order to absorb the wave function at the boundary so as to avoid the reflexion that would otherwise perturb the dynamics. For the decay problems, the absorbing potential simulates the outgoing wave boundary condition, which is imposed when one constructs a Gamow state. Notice that this method is intimately related to the so-called complex absorbing potential (CAP) method, which was developed to compute resonances states in atomic physics [33–35] as well as in nuclear physics [36–38], even though we do not take the limit of vanishing complex absorbing potential.

The paper is organized as follows. In Sec. II, we present the formalism for time-dependent calculations with a complex absorbing potential, which describes quantum mechanically multidimensional tunneling decay problems. In Sec. III, we apply this method to a simple two-dimensional model for spontaneous fission. We compare the results with those in the semiclassical approximation for both the one- and two-dimensional problems. We then summarize the paper in Sec. IV.

II. FORMALISM

To investigate the multidimensional tunneling problem, we solve the TDSE in a finite box. To this end, we add an imaginary potential to the Hamiltonian H , that is, $H' = H + iW(r)$. The imaginary potential $iW(r)$ absorbs the outgoing flux, and should be applied only at the edge of the box in order to not perturb the physical behavior of the decay process. This is effectively equivalent to imposing the outgoing wave boundary condition for resonance states. The TDSE can be integrated as

$$|\Psi(t)\rangle = e^{-\frac{i}{\hbar}t\hat{H}'}|\Psi_0\rangle, \quad (1)$$

where $|\Psi_0\rangle$ is the initial wave function at $t = 0$.

To compute easily the propagator in Eq. (1), we expand the wave function on the biorthogonal basis [39–41] formed by the left and right eigenfunctions of the Hamiltonian H' ,

$$H'|\varphi_i^r\rangle = E_i|\varphi_i^r\rangle \quad \text{and} \quad \langle\varphi_i^l|H' = E_i\langle\varphi_i^l|. \quad (2)$$

Notice that the eigenvalues E_i are complex since the Hamiltonian H' is non-Hermitian. The biorthogonal basis forms the completeness relation as $\sum_i |\varphi_i^r\rangle\langle\varphi_i^l| = \mathbb{1}$, which leads to the

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simple evolution

$$|\Psi(t)\rangle = \sum_i e^{-\frac{i}{\hbar}tE_i} \langle \varphi_i^l | \Psi_0 \rangle |\varphi_i^r\rangle. \quad (3)$$

An advantage of this method is that the evolution of the system can be followed for a very long time. This is particularly suitable for a tunneling process where the lifetime is several orders of magnitudes longer than the characteristic time scale of the system.

Among the eigenstates of H' , we identify the state $|\varphi_i^r\rangle$ which has the smallest value of the imaginary part of the eigenenergy, $E_i = E_i^r - i\Gamma_i/2$, with the physical resonance state. The real part of energy, E_i^r , corresponds to the resonance energy while the imaginary part, Γ_i , corresponds to the resonance width. In fact, it is straightforward with the TDSE to show that this state has the lifetime of $\tau_i = \hbar/\Gamma_i$. In this method, the details of the initial wave function $|\Psi_0\rangle$ is unimportant as long as it has an appreciable overlap with the resonance wave function, $|\varphi_i^r\rangle$.

III. RESULTS

A. Model Hamiltonian

We now apply the formalism presented in the previous section to a fission problem and compare the results with those in the semiclassical approximation. To this end, we employ a two-dimensional fission model considered in Refs. [42,43]. This model consists of the elongation R between the two fission fragments and an intrinsic degree of freedom ξ coupled to the elongation. The Hamiltonian then reads

$$H(R, \xi) = -\frac{\hbar^2}{2M} \frac{\partial^2}{\partial R^2} + U(R) - \frac{\hbar^2}{2m} \frac{\partial^2}{\partial \xi^2} + \frac{1}{2} m \omega^2 \xi^2 + gR\xi. \quad (4)$$

The potential $U(R)$ is chosen to be

$$U(R) = \frac{1}{2} M \Omega^2 R^2 \left(1 - \frac{R}{R_b}\right), \quad (5)$$

in order to form a barrier. The total potential, $V(R, \xi) = U(R) + \frac{1}{2} m \omega^2 \xi^2 + gR\xi$, has a saddle at

$$R_s = \frac{2R_b}{3M\Omega^2} \left(M\Omega^2 - \frac{g^2}{m\omega^2} \right), \quad (6)$$

$$\xi_s = -\frac{2gR_b}{3Mm\Omega^2\omega^2} \left(M\Omega^2 - \frac{g^2}{m\omega^2} \right), \quad (7)$$

with the barrier height of

$$V_b = \frac{1}{6} \left(\frac{2R_b}{3M\Omega^2} \right)^2 \left(M\Omega^2 - \frac{g^2}{m\omega^2} \right)^3. \quad (8)$$

When the intrinsic degree of freedom ξ is neglected, the saddle is at $R_s = 2R_b/3$ with the height of $V_b = 2M\Omega^2 R_b^2/27$ [42].

In the calculations presented below, we use the same parameters as those in Refs. [42,43] except for R_b , which we vary to study the tunneling in the potential with different barrier heights. Those parameters were determined in order to mimic the symmetric fission of ^{234}U with a coupling to the β vibration. The parameters are then taken to be $\hbar\Omega = \hbar\omega = 0.97$ MeV,

$g^2 = Mm\Omega^2/(16\hbar^2)$, $M = 234M_N/4$, and $m = 3AM_N R_0^2/(8\pi)$, where M_N is the nucleon mass, $A = 234$ is the atomic number of the nucleus, and $R_0 = 1.2A^{1/3}$ fm is the equivalent sharp radius (notice that the vertical axis in Fig. 4 in Ref. [43] is actually $R_0\xi$, rather than ξ itself). In the actual calculations, we modify the total potential to a constant value in the outer region to avoid the divergence of the potential, that is, $V(R, \xi) \rightarrow \max(V(R, \xi), -3)$.

B. One-dimensional problem

Before we discuss the tunneling dynamics in the two-dimensional space, let us first solve the problem in one dimension neglecting the ξ degree of freedom. In the one-dimensional problem, the tunneling path is trivial, and the decay lifetime is obtained in the semiclassical approximation as

$$\tau = \frac{2\pi}{\Omega} e^{2S/\hbar}, \quad (9)$$

with

$$S = \int_{R_0}^{R_1} \sqrt{2M[U(R) - E_0]} dR, \quad (10)$$

where E_0 is the energy of the decay state, and R_0 and R_1 are the inner and the outer turning points, respectively, satisfying $U(R_0) = U(R_1) = E_0$. For a cubic potential given by Eq. (5), the semiclassical formula can also be transformed to [20,42]

$$\tau = \frac{1}{\Omega} \sqrt{\frac{\pi\hbar}{60S'}} e^{2S'/\hbar}, \quad (11)$$

with

$$S' = \int_0^{R'_1} \sqrt{2MU(R)} dR, \quad (12)$$

where $U(R'_1) = U(0) = 0$.

To solve the same problem quantum mechanically, we first determine the initial wave function $|\Psi_0\rangle$ by modifying the potential $U(R)$ so that the modified potential has a bound state. To this end, we replace the potential outside the barrier with a constant value [27,44], which we take to be 1 MeV as shown by the dashed line in Fig. 1(a). The corresponding initial wave function is plotted by the dashed line in the lower panel of Fig. 1.

For the complex absorbing potential, we employ the shifted polynomial function [37,38]

$$W(R) = W_0(R - R_a)^2 \theta(R - R_a), \quad (13)$$

with $W_0 = -0.387$ MeV fm $^{-2}$ and $R_a = 14$ fm. We have confirmed that the results do not significantly change even if we vary the value of W_0 and R_a as well as the size of the box.

With this complex potential, we then determine the ensemble of eigenstates and eigenvalues of the Hamiltonian H' defined by Eq. (2). It should be mentioned here that due to the large difference in the order of magnitude between the real and imaginary parts of the resonance energy, it is numerically necessary to use the quadruple precision in the program. With this prescription, physical quantities can be calculated up to about 34 decimal digits. This allows us to calculate a lifetime

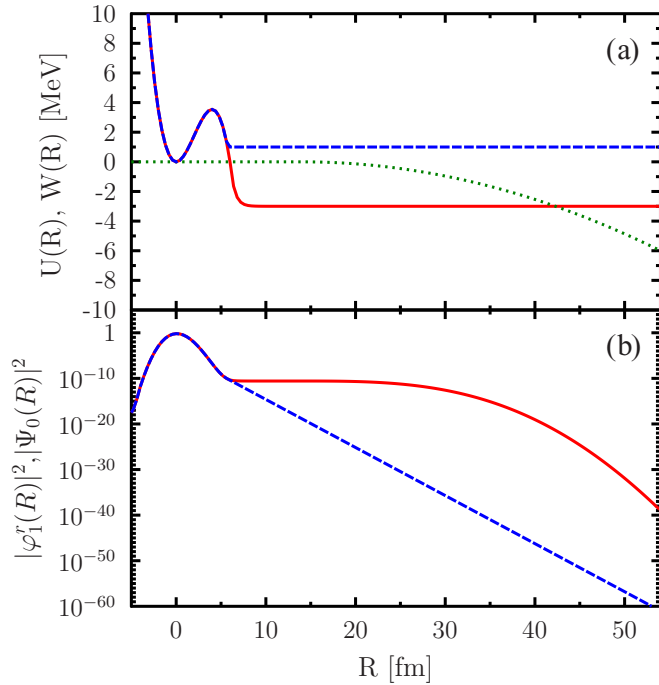


FIG. 1. (Color online) (a) One-dimensional potential $U(R)$ with $R_b = 6$ fm (the red solid line). The figure also shows the modified potential for the initial wave function (the blue dashed line) and the absorbing potential $W(R)$ (the green dotted line). (b) The resultant initial wave function (the blue dashed line) and the resonance wave function (the red solid line).

of the order of 10^9 years when the characteristic time of the system is of the order of 10^{-22} s. Another important parameter in the calculations is the lattice mesh size ΔR , which has to be small enough to describe correctly the tunneling wave function. In our calculations, we take $\Delta R = 0.25$ fm with the finite difference formula with 9 points for the second derivative in the kinetic energy operator.

To select the physical resonance wave function among the eigenstates of H' , we take the lowest energy state $|\varphi_1^r\rangle$ which has the maximum overlap with the initial state. The resulting wave function is shown in Fig. 1(b) by the solid line. We see that the wave function is smoothly damped up to a factor of 10^{30} before reaching the reflecting edge of the lattice at $R = 54$ fm. The decay width can then be read off from the imaginary part of the eigenenergy.

To have a more intuitive picture of the decay process, we compute the time-evolution of the wave function according to Eq. (3). The probability density at different times obtained with a larger value of R_a ($R_a = 54$ fm) is shown in Fig. 2. At $t = 0$, the confining in the modified potential is suddenly removed, and the initial wave function is coupled to the continuum. In the first instant of the dynamics, one can see the emitted wave going outward from the potential barrier. After about $t = 10^4$ fm/c, the emission becomes stationary, and the wave packet has the same shape as the metastable wave function constructed by diagonalizing the Hamiltonian [see the solid line in Fig. 1(b)]. At later time, the wave packet is absorbed exponentially keeping the same spatial shape.

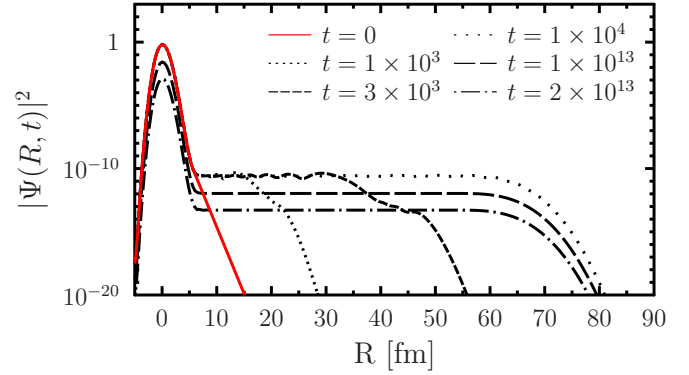


FIG. 2. (Color online) Time evolution of the square of the wave function during the decay process. The time is indicated in unit of fm/c.

The decay width can be calculated with the TDSE by computing the survival probability defined as

$$\mathcal{P}(t) = \int_{-\infty}^{R_{\text{lim}}} |\Psi(R, t)|^2 dR, \quad (14)$$

as a function of time, where R_{lim} is taken outside the barrier. The survival probability obtained with $R_{\text{lim}} = 20$ fm is plotted in Fig. 3 for the potential with $R_b = 11$ fm. We find that both the TDSE method and the imaginary part of the eigenenergy of H' yield $1/\tau = 5.76 \times 10^{-9}$ (1/yr) while the semiclassical approximation yields $1/\tau = 5.69 \times 10^{-9}$ (1/yr). The agreement between the quantal and the semiclassical calculations is rather good for this parameter set, partly because the barrier

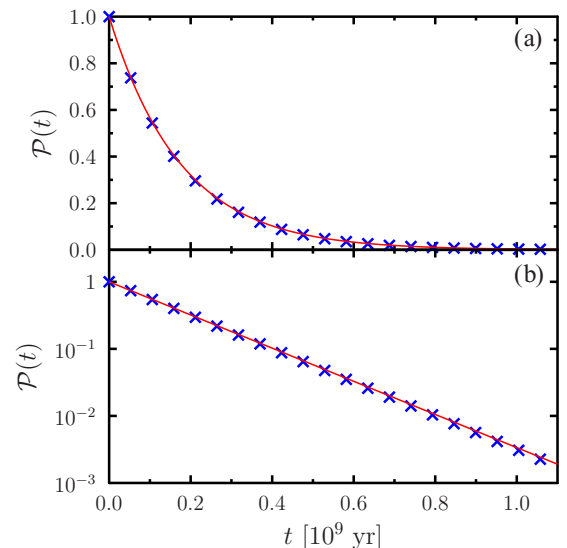


FIG. 3. (Color online) Survival probability as a function of time computed dynamically using the formula (14) with $R_{\text{lim}} = 20$ fm (the blue crosses). The R_b parameter in the potential is taken to be $R_b = 11$ fm. The survival probability is plotted both in the linear scale (the upper panel) and in the logarithmic scale (the lower panel). For a comparison, the figure also shows the survival probabilities for an exponential decay with the decay width obtained from the semiclassical approximation given by Eq. (9) (the red solid line).

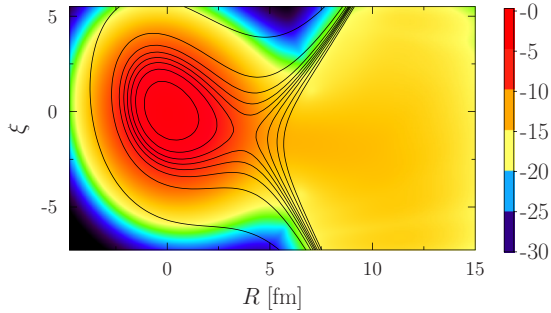


FIG. 4. (Color online) Wave function for the metastable state plotted in the logarithmic scale. This state corresponds to an eigenstate of the Hamiltonian H' for the potential with $R_b = 6$ fm and with the imaginary potential given by Eq. (13) with $W_0 = -0.1$ MeV and $R_a = 11$ fm. The total potential $V(R, \xi)$ is also plotted by the contour lines.

is high and the multiple reflections under the barrier can be neglected.

C. Two-dimensional problem

We now discuss the tunneling dynamics in the two-dimensional surface. To this end, we diagonalize the Hamiltonian H' in two dimensions in a lattice of dimension $R \in [-4.75 \text{ fm} : 16 \text{ fm}]$ and $\xi \in [-7.25 : 5.5]$ with a mesh size of $\Delta R = 0.25$ fm and $\Delta \xi = 0.25$. The imaginary potential is implemented with the same expression as Eq. (13) with $W_0 = -0.1$ MeV and $R_a = 11$ fm. The reference wave function to be used to select the physical resonance state is obtained in a similar manner as in the previous subsection. The wave function for the metastable state is plotted in Fig. 4 for the choice of $R_b = 6$ fm, together with the total potential $V(R, \xi)$ in the contour lines. One can see that this wave function is well confined inside the barrier with a small component outside due to the quantum tunneling, as in the one-dimensional case shown in Fig. 1.

Notice that this wave function corresponds to the wave function after the decay becomes stationary. The pre-stationary decay at the first instant of the decay process is shown in Fig. 5, following the time evolution from the reference wave function as the initial state. One can see that the flow occurs in a small region in the potential energy surface.

We compare the quantum mechanical flow with the semiclassical tunneling path. To this end, we follow the method described in Refs. [20–22]. That is, the semiclassical tunneling path is determined by solving the classical equations of motion in an inverted potential, $V(R, \xi) \rightarrow -V(R, \xi)$. For the initial condition, one can use $R(0) = \epsilon \sqrt{\frac{\hbar}{2M\Omega}} \cos \theta$, $\xi(0) = \epsilon \sqrt{\frac{\hbar}{2m\omega}} \sin \theta$, and $\dot{R}(0) = \dot{\xi}(0) = 0$, where ϵ is a small number. By searching with θ from 0 to 2π , one finds a special angle θ for which the classical path reaches the equipotential surface (and bounces back to the origin if one continues to follow the time evolution of the path), while for other values of θ the classical path is reflected before it reaches the equipotential surface [20–22,43,45]. This special path is referred to as the escape path, and plays an important role in the semiclassical

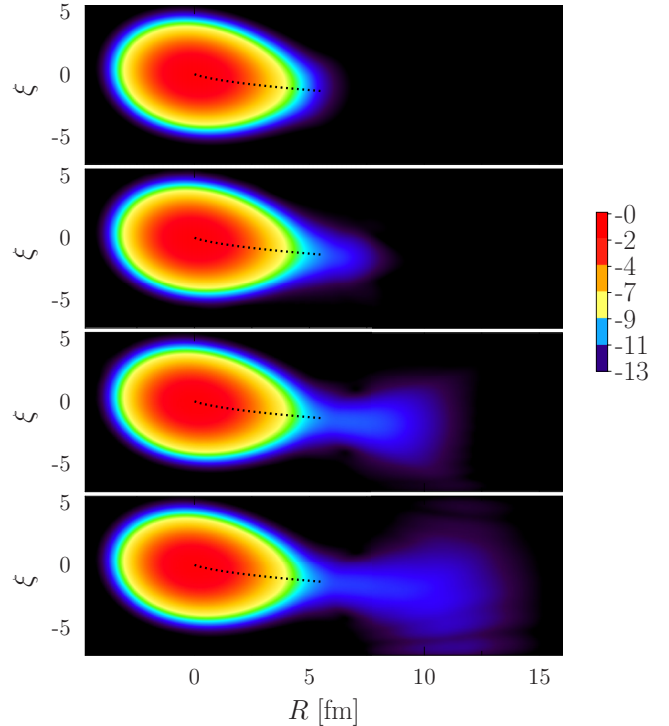


FIG. 5. (Color online) Time evolution of the two-dimensional wave function at time $t = 0, 250, 500,$ and 1250 fm/c. For a comparison, the semiclassical tunneling path is also shown with the dashed line.

theory of multidimensional quantum tunneling. The escape path so obtained is denoted by the dashed line in Fig. 5 (there is only one escape path for the potential considered in this paper). It is remarkable that the quantum mechanical time evolution almost follows the semiclassical path. We have confirmed that the tunneling path obtained by minimizing the classical action with the algorithm in Ref. [46] provides the same result.

To make a further comparison of the tunneling path, we compute the flux, $\mathbf{j} = (j_R, j_\xi)$, from the resonance wave

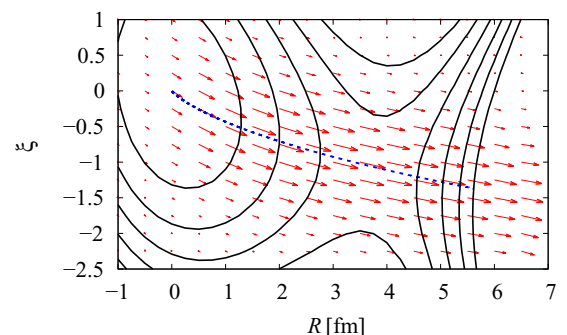


FIG. 6. (Color online) Comparison of the quantum mechanical flux computed with Eqs. (15) and (16) (the red arrows) with the semiclassical escape path (the blue dashed line). The two-dimensional potential energy surface is also shown by the black contour lines.

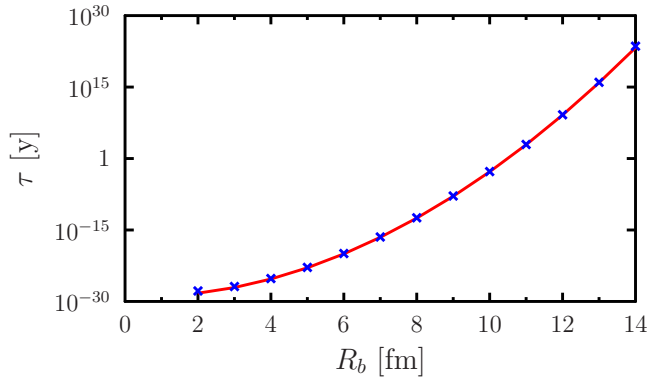


FIG. 7. (Color online) Fission lifetime as a function of the R_b parameter in the potential $U(R)$. The lifetime computed quantum mechanically is denoted by the blue crosses, while that evaluated with the semiclassical method using Eq. (17) is denoted by the red line.

function as

$$j_R = \frac{\hbar}{2iM} \left(\varphi_1^{r*} \frac{\partial \varphi_1^r}{\partial R} - \varphi_1^r \frac{\partial \varphi_1^{r*}}{\partial R} \right), \quad (15)$$

and

$$j_\xi = \frac{\hbar}{2im} \left(\varphi_1^{r*} \frac{\partial \varphi_1^r}{\partial \xi} - \varphi_1^r \frac{\partial \varphi_1^{r*}}{\partial \xi} \right). \quad (16)$$

The flux is shown in Fig. 6 and is compared to the semiclassical escape path. The quantum flow is a collection of all the trajectories connecting the region around the origin with the continuum region. We see that the main quantum mechanical flow is systematically parallel to the semiclassical trajectory. The semiclassical path can thus be regarded as the mean trajectory of the quantum flow, whereas the quantum flow takes into account the quantum mechanical fluctuation around the classical trajectory.

In addition to the agreement for the tunneling path, we also compare in Fig. 7 the resulting fission lifetime. Varying the R_b parameter in the potential $U(R)$, we obtain a range of fission lifetimes from the characteristic time of the system to a lifetime longer than the actual lifetime of the ^{238}U nucleus. The quantum mechanical lifetime is computed from the imaginary part of the resonance energy, while the lifetime in the semiclassical approximation is evaluated using the Eq. (11) with the action integral evaluated along the escape

path \mathcal{P} in the imaginary time,

$$S' = \int_{\mathcal{P}} (M\dot{R}^2 + m\dot{\xi}^2) d\tau. \quad (17)$$

The latter is equivalent to approximating the pre-exponential factor by that for a one-dimensional problem with a cubic potential along the escape path [43]. In the range shown in the figure, a good agreement is found between the two methods, the maximum deviation being up to about 20%. Evidently, the semiclassical method provides a good approximation to the multidimensional tunneling problem for this Hamiltonian.

IV. SUMMARY

We have presented a full-quantum method to study a decay lifetime as well as the tunneling dynamics. To this end, we have introduced a complex absorbing potential, which is effectively equivalent to the outgoing boundary condition, and constructed the biorthogonal basis. We have shown that this method provides a good tool to follow the time evolution of a system over a very long time. This enables one to compute the decay lifetime from a few fm/c to the order of 10^9 years. A comparison with the semiclassical approximation has shown a good agreement between the two methods. It has been shown that the average of fissioning flux in a multidimensional plane corresponds to the semiclassical tunneling path. For the decay lifetime, the two methods yield similar values to one another, where the maximum difference is only about 20%.

The quantum mechanical method discussed in this paper provides a good alternative to the semiclassical method. It provides an intuitive picture of multidimensional quantum tunneling, including the quantum mechanical fluctuation of the classical path. It will also provide a convenient method when a bifurcation of the tunneling path is important, e.g., in the presence of a competition of several fission modes. We plan to apply this method, both for dynamics and a determination of resonance state, to more realistic systems using the constrained mean-field theory for the potential energy surface and/or the time-dependent generator coordinate method [13,14,47]. We will report it in a separate publication.

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- [1] M. Brack, J. Damgaard, A. J. Jensen, H. C. Pauli, V. M. Strutinsky, and C. Y. Wong, *Rev. Mod. Phys.* **44**, 320 (1972).
- [2] P. Möller, D. G. Madland, A. J. Sierk, and A. Iwamoto, *Nature* **409**, 785 (2001).
- [3] P. Möller, A. J. Sierk, T. Ichikawa, A. Iwamoto, R. Bengtsson, H. Uhrenholt, and S. Aberg, *Phys. Rev. C* **79**, 064304 (2009).
- [4] J. Skalski, *Phys. Rev. C* **77**, 064610 (2008).

- [5] A. Staszczak, A. Baran, J. Dobaczewski, and W. Nazarewicz, *Phys. Rev. C* **80**, 014309 (2009).
- [6] A. Baran, J. A. Sheikh, J. Dobaczewski, W. Nazarewicz, and A. Staszczak, *Phys. Rev. C* **84**, 054321 (2011).
- [7] J. Sadhukhan, K. Mazurek, A. Baran, J. Dobaczewski, W. Nazarewicz, and J. A. Sheikh, *Phys. Rev. C* **88**, 064314 (2013).
- [8] J. D. McDonnell, W. Nazarewicz, J. A. Sheikh, A. Staszczak, and M. Warda, *Phys. Rev. C* **90**, 021302(R) (2014).

- [9] J. Sadhukhan, J. Dobaczewski, W. Nazarewicz, J. A. Sheikh, and A. Baran, *Phys. Rev. C* **90**, 061304(R) (2014).
- [10] M. Warda and J. L. Egidio, *Phys. Rev. C* **86**, 014322 (2012).
- [11] R. Rodriguez-Guzman and L. M. Robledo, *Phys. Rev. C* **89**, 054310 (2014).
- [12] S. A. Giuliani, L. M. Robledo, and R. Rodriguez-Guzman, *Phys. Rev. C* **90**, 054311 (2014).
- [13] H. Goutte, J. F. Berger, P. Casoli, and D. Gogny, *Phys. Rev. C* **71**, 024316 (2005).
- [14] R. Bernard, H. Goutte, D. Gogny, and W. Younes, *Phys. Rev. C* **84**, 044308 (2011).
- [15] H. Abusara, A. V. Afanasjev, and P. Ring, *Phys. Rev. C* **82**, 044303 (2010); **85**, 024314 (2012).
- [16] B.-N. Lu, E.-G. Zhao, and S.-G. Zhou, *Phys. Rev. C* **85**, 011301 (2012); B.-N. Lu, J. Zhao, E.-G. Zhao, and S.-G. Zhou, *ibid.* **89**, 014323 (2014).
- [17] J. Zhao, B.-N. Lu, D. Vretenar, E.-G. Zhao, and S.-G. Zhou, *Phys. Rev. C* **91**, 014321 (2015).
- [18] T. Bürvenich, M. Bender, J. A. Maruhn, and P.-G. Reinhard, *Phys. Rev. C* **69**, 014307 (2004).
- [19] W. D. Myers and W. J. Swiatecki, *Nucl. Phys. A* **601**, 141 (1996).
- [20] A. Schmid, *Ann. Phys. (NY)* **170**, 333 (1986).
- [21] T. Kindo and A. Iwamoto, *Phys. Lett. B* **225**, 203 (1989).
- [22] A. Iwamoto, *Z. Phys. A* **349**, 265 (1994).
- [23] N. Carjan, O. Serot, and D. Strottman, *Z. Phys. A* **349**, 353 (1994).
- [24] P. Talou, D. Strottman, and N. Carjan, *Phys. Rev. C* **60**, 054318 (1999).
- [25] D. Lacroix, J. A. Scarpaci, and Ph. Chomaz, *Nucl. Phys. A* **658**, 3 (1999).
- [26] T. Maruyama, T. Oishi, K. Hagino, and H. Sagawa, *Phys. Rev. C* **86**, 044301 (2012).
- [27] T. Oishi, K. Hagino, and H. Sagawa, *Phys. Rev. C* **90**, 034303 (2014).
- [28] M. Ueda, K. Yabana, and T. Nakatsukasa, *Phys. Rev. C* **67**, 014606 (2003).
- [29] T. Nakatsukasa and K. Yabana, *Phys. Rev. C* **71**, 024301 (2005).
- [30] P.-G. Reinhard, P. D. Stevenson, D. Almeded, J. A. Maruhn, and M. R. Strayer, *Phys. Rev. E* **73**, 036709 (2006).
- [31] G. Scamps, D. Lacroix, G. F. Bertsch, and K. Washiyama, *Phys. Rev. C* **85**, 034328 (2012).
- [32] J. E. Elenewski and H. Chen, *Phys. Rev. B* **90**, 085104 (2014).
- [33] U. V. Riss and H.-D. Meyer, *J. Phys. B* **26**, 4503 (1993).
- [34] S. Feuerbacher, T. Sommerfeld, R. Santra, and L. S. Cederbaum, *J. Chem. Phys.* **118**, 6188 (2003).
- [35] R. Santra and L. S. Cederbaum, *J. Chem. Phys.* **115**, 6853 (2001).
- [36] H. Masui and Y. K. Ho, *Phys. Rev. C* **65**, 054305 (2002).
- [37] M. Ito and K. Yabana, *Prog. Theor. Phys.* **113**, 1047 (2005).
- [38] R. Otani, R. Kageyama, M. Iwasaki, M. Kudo, M. Tomita, and M. Ito, *Phys. Rev. C* **90**, 034316 (2014).
- [39] P. Morse and H. Feshbach, *Methods of Theoretical Physics* (McGraw-Hill, New York, 1952), p. 884.
- [40] M. S. Hussein, M. P. Pato, and A. F. R. de Toledo Piza, *Phys. Rev. C* **51**, 846 (1995).
- [41] K. Hagino and N. Takigawa, *Phys. Rev. C* **58**, 2872 (1998).
- [42] D. M. Brink, M. C. Nemes, and D. Vautherin, *Ann. Phys. (NY)* **147**, 171 (1983).
- [43] N. Takigawa, K. Hagino, and M. Abe, *Phys. Rev. C* **51**, 187 (1995).
- [44] S. A. Gurvitz, P. B. Semmes, W. Nazarewicz, and T. Vertse, *Phys. Rev. A* **69**, 042705 (2004).
- [45] D. M. Brink, in *The Proceedings of the Peierls 80th Birthday Symposium, A Breadth of Physics*, edited by R. H. Dalitz and R. B. Stinchcombe (World Scientific, Singapore, 1988), p. 109.
- [46] A. Baran and K. Pomorski, *Nucl. Phys. A* **361**, 83 (1981).
- [47] P.-G. Reinhard, R. Y. Cusson, and K. Goeke, *Nucl. Phys. A* **398**, 141 (1983).