

Low-lying collective excited states in non-integrable system based on stationary phase approximation to the path integral

Fang Ni¹ and Takashi Nakatsukasa^{2, 1, 3}

¹*Faculty of Pure and Applied Sciences, University of Tsukuba, Tsukuba 305-8571, Japan*

²*Center for Computational Sciences, University of Tsukuba, Tsukuba 305-8577, Japan*

³*iTHES Research Group, RIKEN, Wako 351-0198, Japan*

Background: This part would describe the context needed to understand what the paper is about.

Purpose: This part would state the purpose of the present paper.

Method: This part describe the methods used in the paper.

Results: This part would summarize the results.

Conclusions: This part would state the conclusions of the paper.

I. INTRODUCTION

II. THEORETICAL FRAMEWORK

A. One-dimensional adiabatic self-consistent collective coordinate method

We review the one-dimensional adiabatic self-consistent collective coordinate method (ASCC) briefly. The classical Hamilton's equation which is identical to the TDHFB equation is described by canonical variables $\{\xi^\alpha, \pi_\alpha\}$. If we assume the collective motion is slow motion, it allows us to expand momenta π up to second order. The multi-dimensional Hamiltonian is

$$\mathcal{H} = V(\xi) + \frac{1}{2} B^{\alpha\beta}(\xi) \pi_\alpha \pi_\beta \quad (\text{II.1})$$

with potential $V(\xi)$ and reciprocal mass parameter $B^{\alpha\beta}(\xi)$ defined by

$$V(\xi) = \mathcal{H}(\xi, \pi = 0) \quad (\text{II.2})$$

$$B^{\alpha\beta}(\xi) = \left. \frac{\partial^2 \mathcal{H}(\xi, \pi)}{\partial \pi_\alpha \partial \pi_\beta} \right|_{\pi=0}. \quad (\text{II.3})$$

If there is a one-dimensional collective motion which is decoupled from other intrinsic motions, the collective motion can be described by a set of canonical variable $\{q, p\}$, the one dimensional collective Hamiltonian is

$$\mathcal{H}_{coll} = \bar{V}(q) + \frac{1}{2} \bar{B}^{-1}(q) p^2. \quad (\text{II.4})$$

To obtain the collective Hamiltonian, we need to consider extended adiabatic transformation

$$q = q^1 = f^1(\xi) + \frac{1}{2} f^{(1)1\alpha\beta}(\xi) \pi_\alpha \pi_\beta \quad (\text{II.5})$$

$$\xi^\alpha = g^\alpha(q) + \frac{1}{2} g^{(1)\alpha 11}(\xi) p_1 p_1 \quad (\text{II.6})$$

and

$$p_1 = g_{,1}^\alpha \pi_\alpha \quad (\text{II.7})$$

$$\pi_\alpha = f_{,\alpha}^1 p_1, \quad (\text{II.8})$$

where the index 1 indicates the collective degree of freedom, and the comma indicates the partial derivative ($f_{,\alpha}^1 = \partial f^1 / \partial \xi^\alpha$). With these relations, collective potential $\bar{V}(q)$ and collective mass parameter $\bar{B}(q)$ can be transformed by

$$\bar{V}(q) = V(\xi) \quad (\text{II.9})$$

$$\bar{B}^{-1}(q) = f_{,\alpha}^1 B^{\alpha\beta}(\xi) f_{,\beta}^1, \quad (\text{II.10})$$

Before showing the ASCC basic equations, we consider the constants of motion in TDHFB dynamics. Since nuclei is self-bound systems without external potential, the ground state obtained from mean-field theory violates the symmetry (e.g. deformation, pairing). In such system, the constants of motion corresponding the Nambu-Goldstone (NG) mode emerges in TDHFB dynamics. Therefore, we are mostly interested in the systems with collective motion separated from NG mode.

We know that the most of conserved quantities, such as total angular momentum J and total particle number N , are not only expressed by one-body Hermitian operators, but also has real matrix elements in the quasi-particle basis. Such classical variable \mathcal{P} corresponding to the conserved quantity can be expanded as

$$\mathcal{P}(\xi, \pi) = f^I(\xi) + \frac{1}{2} f^{(1)I\alpha\beta}(\xi) \pi_\alpha \pi_\beta. \quad (\text{II.11})$$

The index I indicates the degree of freedom for NG mode. Since the conserved quantity must fulfill $\{\mathcal{P}, \mathcal{H}\}_{PB} = 0$, it leads

$$f_{,\alpha}^I B^{\alpha\beta} - f^{(1)I\alpha\beta} V_{,\alpha} = 0. \quad (\text{II.12})$$

Including the NG mode, $f^1(\xi)$ and $g^\alpha(q)$ are obtained by solving ASCC basic equations

$$\delta H_M(\xi, \pi) = 0 \quad (\text{II.13})$$

$$\tilde{B}^{\beta\gamma} V_{;\gamma\alpha} f_{,\beta}^1 = \omega^2 f_{,\alpha}^1, \quad \tilde{B}^{\beta\gamma} V_{;\gamma\alpha} g_{,1}^\alpha = \omega^2 g_{,1}^\beta. \quad (\text{II.14})$$

The first equation (II.13) is called moving-frame HFB equation. The moving-frame Hamiltonian \mathcal{H}_M with constraints on collective coordinate and constant of motion is

$$\mathcal{H}_M(\xi, \pi) = \mathcal{H}(\xi, \pi) - \lambda_1 q^1(\xi, \pi) - \lambda_I \mathcal{P}(\xi, \pi). \quad (\text{II.15})$$

The second equation (II.14) is called moving-frame QRPA equation. $\tilde{B}^{\alpha\beta}$ is the modified mass parameter

$$\begin{aligned}\tilde{B}^{\alpha\beta}(\xi) &= \frac{\partial^2 \mathcal{H}_M}{\partial \pi_\alpha \partial \pi_\beta} \\ &= B^{\alpha\beta}(\xi) - \lambda_I f^{(1)I\alpha\beta}(\xi) - \lambda_1 f^{(1)1\alpha\beta}(\xi).\end{aligned}\quad (\text{II.16})$$

and the covariant derivative $V_{;\gamma\alpha}$ is defined by

$$V_{;\alpha\beta} = V_{,\alpha\beta} - \Gamma_{\alpha\beta}^\gamma V_{,\gamma}, \quad (\text{II.17})$$

where the affine connection is $\Gamma_{\beta\gamma}^\alpha = \frac{1}{2}B^{\alpha\delta}(B_{\delta\beta,\gamma} + B_{\delta\gamma,\beta} - B_{\beta\gamma,\delta})$. If we assume that the collective coordinate q is geodesic, $\tilde{B}^{\beta\gamma}V_{;\gamma\alpha}$ can be simplified after several steps

$$\tilde{B}^{\alpha\gamma}V_{;\gamma\beta} = \tilde{B}^{\alpha\gamma}(V_{,\gamma\beta} - \lambda_I f_{,\gamma\beta}^I) + \frac{1}{2}\tilde{B}_{,\beta}^{\alpha\gamma}V_{,\gamma}. \quad (\text{II.18})$$

We discuss the practical solution to derive the one-dimensional collective coordinate. We neglect $f^{(1)1\alpha\beta}(\xi)$ because it is supposed to be negligibly small in the previous study. On the other hand, from (II.12) and (II.14), $f^{(1)I\alpha\beta}(\xi)$ is necessary information to promise $\tilde{B}^{\beta\gamma}V_{;\gamma\alpha}f_{,\beta}^I = 0$, which means zero mode. In most of case, we know the explicit form of $\mathcal{P}(\xi, \pi)$. The procedures to obtain the collective path are (see Figure ??):

1. Find the energy minimum point in energy surface by solving (II.13) (normal HFB equation). We usually set $q^1 = 0$ at the energy minimum point.
2. Using (II.18), diagonalize (II.14) and choose the lowest mode basically*.
3. The right eigenvector $g_{,1}^\alpha$ tell us the direction which system moves in energy surface. Use the relation $d\xi^\alpha = g_{,1}^\alpha dq^1$ to decide the neighborhood point $q^1 \rightarrow q^1 + dq^1$ in the collective path.
4. Find the non-equilibrium energy minimum point by solving (II.13) at the neighborhood point, and obtain the chemical potential $\lambda_I(q)$.
5. Iterate (2)~(4) under the same direction in energy surface.
6. Do (2)~(5) for the opposite direction of the collective path in energy surface.

The scale of the collective coordinate is not unique because of the uncertainty of the eigenvectors in (II.14). To unify the scale, the simplest procedure is to keep the collective mass parameter $\bar{B}^{-1}(q) = 1$ by renormalizing eigenvectors in (II.10).

* When eigenvalues cross on the collective path, the choice of adiabatic path or diabatic path is a critical problem

B. Stationary-phase approximation to the path integral

In integrable system, we can apply the stationary phase approximation to the path integral (SPA) to obtain collective excited states. The concept of SPA is introduced in the previous study. Because the one-dimensional collective path extracted from TDHFB degrees of freedom is integrable system, SPA is supposed to be available for non-integrable system via ASCC.

Based on SPA, the k -th excited state $|\psi_k\rangle$ can be constructed by a periodic TDHFB trajectory

$$|\psi_k\rangle = \oint d\mu(Z^{(k)}) |Z^{(k)}\rangle e^{i\mathcal{T}[Z^{(k)}]/\hbar}. \quad (\text{II.19})$$

The integration indicates we integrate a periodic TDHFB trajectory for one period in TDHFB phase space. The invariant measure $d\mu(Z)$ is defined by the unity condition $\int d\mu(Z) |Z\rangle \langle Z| = 1$. The action integral \mathcal{T} is defined by

$$\mathcal{T}[Z] \equiv \int_0^t \langle Z(t') | i\hbar \frac{\partial}{\partial t'} | Z(t') \rangle dt'. \quad (\text{II.20})$$

We try to combine SPA with ASCC. The key point is to find the correspondence between TDHFB phase space and collective subspace in a TDHFB trajectory (See Fig. 1). We consider the time-dependent state vector $|Z(t)\rangle$ is in a collective subspace. With pairing correlation, $|Z(t)\rangle$ can be expressed as

$$|Z(t)\rangle = |\Phi, J; q, p\rangle = e^{-i\Phi\hat{J}} |J; q, p\rangle, \quad (\text{II.21})$$

where Φ is the total gauge angle and $J = N/2$ is the conjugate momentum corresponding total particle number. The second equal sign indicates an intrinsic state $|J; q, p\rangle$ rotates in the gauge space. Since $[H, J] = 0$, the classical Hamiltonian becomes

$$\begin{aligned}\mathcal{H} &= \langle \Phi, J; q, p | H | \Phi, J; q, p \rangle \\ &= \langle J; q, p | H | J; q, p \rangle \equiv \mathcal{H}_J(q, p)\end{aligned}\quad (\text{II.22})$$

has no dependence of Φ . Therefore, the classical Hamiltonian is the function of (q, p) in a fixed particle number system, and correspond to (II.4). From ASCC, we can obtain the static state vector at each point of q , namely $|J; q, p = 0\rangle$. To construct $|J; q, p\rangle$, we use (II.8) at (q, p) which we want to know in the collective phase space. Usually, we are interested in the points $(q^{(k)}, p^{(k)})$ at the k -th TDHFB trajectory. Such k -th TDHFB trajectory is obtained from EBK quantization condition (k : integer)

$$\begin{aligned}\mathcal{T}_0[Z^{(k)}] &= \oint \langle Z^{(k)}(t') | i\hbar \frac{\partial}{\partial t'} | Z^{(k)}(t') \rangle dt' \\ &= \oint p^{(k)} dq^{(k)} = 2k\pi.\end{aligned}\quad (\text{II.23})$$

The procedures to obtain excited states from SPA via ASCC are:

1. Obtain the one-dimensional collective coordinate from ASCC.

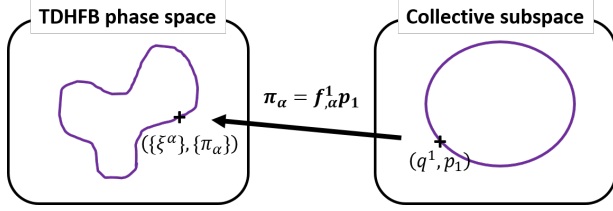


FIG. 1: For a TDHFB trajectory, the correspondence between TDHFB phase space and collective subspace.

2. Using collective Hamiltonian, obtain k -th excitation energy and TDHFB trajectory under EBK quantization condition (II.23).
3. For k -th TDHFB trajectory, construct the state vector $|J; q, p\rangle$ by using (II.8).
4. Calculate action integral (III.22) in k -th TDHFB trajectory.
5. Using (II.19), obtain k -th excited state.

III. APPLICATION IN PAIRING MODEL

We study the low-lying excited 0^+ states in multi-level pairing model by applying SPA via ASCC. The Hamiltonian of the pairing model is given in terms of single-particle energies ϵ_l and the pairing strength g as

$$\begin{aligned} H &= \sum_l \epsilon_l n_l - g \sum_{l,l'} S_l^+ S_{l'}^- \\ &= \sum_l \epsilon_l (2S_l^0 + \Omega_l) - g S^+ S^-, \end{aligned} \quad (\text{III.1})$$

where we use the SU(2) quasi-spin operators, $\mathbf{S} = \sum_l \mathbf{S}_l$, with

$$S_l^0 = \frac{1}{2} (\sum_m a_{lm}^\dagger a_{lm} - \Omega_l), \quad (\text{III.2})$$

$$S_l^+ = \sum_{m>0} a_{lm}^\dagger a_{l\bar{m}}^\dagger, \quad S_l^- = S_l^{+\dagger}. \quad (\text{III.3})$$

Each single-particle energy ϵ_l possesses $(2\Omega_l)$ -fold degeneracy ($\Omega_l = j_l + 1/2$) and $\sum_{m>0}$ indicates the summation over $m = 1/2, 3/2, \dots$, and $\Omega_l - 1/2$. The occupation number of each level l is given by $n_l = \sum_m a_{lm}^\dagger a_{lm} = 2S_l^0 + \Omega_l$. The quasi-spin operators satisfy the commutation relations

$$[S_l^0, S_{l'}^\pm] = \pm \delta_{ll'} S_l^\pm, \quad [S_l^+, S_{l'}^-] = 2\delta_{ll'} S_l^0. \quad (\text{III.4})$$

The magnitude of quasi-spin for each level is $S_l = \frac{1}{2}(\Omega_l - \nu_l)$, where ν_l is the seniority quantum number, namely the number of unpaired particle at each level l . In the present study, we only consider seniority zero states with $\nu = \sum_l \nu_l = 0$. The residual two-body interaction only consists of monopole pairing interaction which couples

two particles to zero angular momentum. We obtain exact solutions either by solving Richardson equation [?] or by diagonalizing the Hamiltonian using the quasi-spin symmetry.

A. Classical form of TDHFB Hamiltonian

The coherent state for the seniority $\nu = 0$ states ($S_l = \Omega_l/2$) is constructed as

$$|Z(t)\rangle = \prod_l (1 + |Z_l(t)|^2)^{-\Omega_l/2} \exp[Z_l(t) S_l^+] |0\rangle \quad (\text{III.5})$$

where $|0\rangle$ is the vacuum (zero particle) state, $Z_l(t)$ are time-dependent complex variables which describe motion of the system. In the SU(2) quasi-spin representation, $|0\rangle = \prod_l |S_l, -S_l\rangle$. The coherent state $|Z(t)\rangle$ is a superposition of states with different particle numbers without unpaired particles. In the present pairing model, the coherent state is the same as the time-dependent BCS wave function with $Z_l(t) = v_l(t)/u_l(t)$, where $(u_l(t), v_l(t))$ are the time-dependent BCS u, v factors.

The TDHFB equation can be derived from the time-dependent variational principle (we set $\hbar = 1$),

$$\delta S = 0, \quad S \equiv \int \mathcal{L}(t) dt = \int \langle Z(t) | i \frac{\partial}{\partial t} - H | Z(t) \rangle dt. \quad (\text{III.6})$$

After transformation of the complex variables $Z_l = \tan \frac{\theta_l}{2} e^{-i\chi_l}$ ($0 \leq \theta \leq \pi$) and several steps of the derivation, the Lagrangian \mathcal{L} and the expectation value of Hamiltonian become

$$\mathcal{L}(t) = \sum_l \frac{\Omega_l}{2} (1 - \cos \theta_l) \dot{\chi}_l - \mathcal{H}(Z, Z^*), \quad (\text{III.7})$$

$$\begin{aligned} \mathcal{H}(Z, Z^*) &\equiv \langle Z | H | Z \rangle \\ &= \sum_l \epsilon_l \Omega_l (1 - \cos \theta_l) - \frac{g}{4} \sum_l \Omega_l [\Omega_l (1 - \cos^2 \theta_l) + (1 - \cos \theta_l)^2] \\ &\quad - \frac{g}{4} \sum_{l_1 \neq l_2} \Omega_{l_1} \Omega_{l_2} \sqrt{(1 - \cos^2 \theta_{l_1})(1 - \cos^2 \theta_{l_2})} e^{-i(\chi_{l_1} - \chi_{l_2})}. \end{aligned} \quad (\text{III.8})$$

Here, we choose χ_l as canonical coordinates. Their conjugate momenta are given by

$$j_l \equiv \frac{\partial \mathcal{L}}{\partial \dot{\chi}_l} = \frac{\Omega_l}{2} (1 - \cos \theta_l). \quad (\text{III.9})$$

χ_l represent a kind of gauge angle of each level, and j_l are related to the occupation number of each level, $2j_l = n_l$. Therefore, the TDHFB Hamiltonian can be represented by canonical variables $\mathcal{H}(Z, Z^*) = \mathcal{H}(\{\chi_l\}, \{j_l\})$, and the TDHFB equation is equivalent to the Hamilton's equation

$$\dot{\chi}_l = \frac{\partial \mathcal{H}}{\partial j_l}, \quad \dot{j}_l = -\frac{\partial \mathcal{H}}{\partial \chi_l}. \quad (\text{III.10})$$

B. Application into ASCC

We construct 1D collective subspace from ASCC theory. Because the first order with respect to χ_l is zero in TDHFB Hamiltonian, we assume j_l as coordinates and χ_l as conjugate momenta. The canonical variables are $(j_l, -\chi_l)$ and fulfill $\{j_l, -\chi_l'\}_{PB} = \delta_{ll'}$. We expand the classical Hamiltonian up to second order with respect to χ_l . The classical Hamiltonian becomes

$$\mathcal{H}(\{\chi_l\}, \{\pi_l\}) \approx V(j) + \frac{1}{2} B^{\alpha\beta}(j) \chi_\alpha \chi_\beta, \quad (\text{III.11})$$

where potential $V(j)$ and reciprocal mass parameter $B^{\alpha\beta}(j)$ are defined as

$$\begin{aligned} V(j) &= \mathcal{H}(\chi = 0, j) \\ &= \sum_l 2\epsilon_l j_l - g \sum_l \left(\Omega_l j_l - j_l^2 + \frac{j_l^2}{\Omega_l} \right) \\ &\quad - g \sum_{l_1 \neq l_2} \sqrt{j_{l_1} j_{l_2} (\Omega_{l_1} - j_{l_1}) (\Omega_{l_2} - j_{l_2})} \end{aligned} \quad (\text{III.12})$$

$$\begin{aligned} B^{\alpha\beta}(j) &= \left. \frac{\partial^2 \mathcal{H}}{\partial \chi_\alpha \partial \chi_\beta} \right|_{\chi=0} \\ &= \begin{cases} 2g \sum_{l \neq \alpha} \sqrt{j_l j_\alpha (\Omega_l - j_l) (\Omega_\alpha - j_\alpha)} & (\alpha = \beta) \\ -2g \sqrt{j_\alpha j_\beta (\Omega_\alpha - j_\alpha) (\Omega_\beta - j_\beta)} & (\alpha \neq \beta) \end{cases} \end{aligned} \quad (\text{III.13})$$

Using the TDHFB Hamiltonian, we can apply ASCC explained in Sec. II A by replacing $\xi \rightarrow j$ and $\pi \rightarrow -\chi$.

We consider the treatment for the constant of motion in pairing model. In the TDHFB Hamiltonian, there is no dependence about the total gauge angle

$$\Phi = \frac{1}{L} \sum_l \chi_l. \quad (\text{III.14})$$

It indicates the conjugate momentum

$$J \equiv \frac{\partial \mathcal{L}}{\partial \dot{\Phi}} = \sum_l j_l = \frac{N}{2} \quad (\text{III.15})$$

is conserved quantity. From this equation, we can find J is not dependent of χ , which means $f^{(1)I\alpha\beta}$ in (II.11) is zero. Therefore, $B^{\beta\gamma} V_{;\gamma\alpha} f_{;\beta}^I = 0$ is promised exactly. Also, the second derivative of J with respect to j vanishes, which indicates $f_{;\gamma\beta}^I$ in (II.18) is zero. It indicates the chemical potential $\lambda_I(q)$ is not necessary information to calculate the matrix element in moving-frame QRPA equation.

With the properties of the constant of motion, we discuss the practical solution of ASCC in pairing model. If we ignore the higher order term $f^{(1)1\alpha\beta}$, we can practice local harmonic equation (LHE)

$$B^{\beta\gamma} V_{;\gamma\alpha} f_{;\beta}^1 = \omega^2 f_{;\alpha}^1, \quad B^{\beta\gamma} V_{;\gamma\alpha} g_{;1}^\alpha = \omega^2 g_{;1}^\beta, \quad (\text{III.16})$$

which simplifies the moving-frame QRPA equation (II.14) by replacing $\tilde{B}^{\beta\gamma}$ into $B^{\beta\gamma}$. Under (III.16), the

eigenvalue of spurious mode corresponding pairing rotation is exact zero. To guarantee the state vectors $|Z(q)\rangle$ are in the same total gauge angle $\Phi(q) = -\frac{1}{L} p_1 \sum_l f_{;l}^1 = 0$, we correct the eigenvectors by

$$f_{;\alpha}^1 \rightarrow f_{;\alpha}^1 - \frac{1}{L} \sum_l f_{;l}^1 \quad (\text{III.17})$$

after solving (III.16) at each point of q .

C. Application into SPA

We derive the explicit form of excited states in (II.19). From (II.21) and (III.15), the state vector $|\Phi, J; q, p\rangle$ becomes

$$|\Phi, J; q, p\rangle = \sum_{\{j_l\}} e^{-i\Phi \sum_l j_l} |J; q, p\rangle. \quad (\text{III.18})$$

Here, $|J; q, p\rangle$ can be expanded in SU(2) quasispin basis

$$|J; q, p\rangle = \sum_{\{j_l\}} A_{j_l}(q, p) |\cdots; S_l, -S_l + k_l, \cdots\rangle, \quad (\text{III.19})$$

where

$$\begin{aligned} A_{j_l}(q, p) &= \prod_l \left(\frac{1 - \cos \theta_l}{2} \right)^{j_l/2} \left(\frac{1 + \cos \theta_l}{2} \right)^{(\Omega_l - j_l)/2} \\ &\quad \times \sqrt{\frac{(\Omega_l)!}{j_l! (\Omega_l - j_l)!}} e^{-ij_l \chi_l} \end{aligned} \quad (\text{III.20})$$

is in fixed gauge angle ($\Phi = 0$).

Due to J is conserved quantity, the action integral can be divided into two terms

$$\mathcal{T}(\Phi, J; q, p) = \mathcal{T}_{\text{intr}}(t) + J\Phi, \quad (\text{III.21})$$

where the intrinsic action integral $\mathcal{T}_{\text{intr}}(t)$ becomes

$$\begin{aligned} \mathcal{T}_{\text{intr}}(t) &= \int \langle J; q, p | i \frac{\partial}{\partial t} | J; q, p \rangle dt \\ &= \int \sum_l j_l d\chi_l. \end{aligned} \quad (\text{III.22})$$

The important point is $\mathcal{T}_{\text{intr}}(t) \neq \int p dq$ at each t . For closed TDHFB trajectory, only $\mathcal{T}_{\text{intr}}(t)$ contributes to the excited states. In SU(2) representation, the invariant measure is

$$\begin{aligned} d\mu(Z) &= \prod_l \frac{\Omega_l + 1}{\pi} (1 + |Z_l|^2)^{-2} d\text{Re} Z d\text{Im} Z \\ &= \prod_l \frac{\Omega_l + 1}{4\pi} d\cos \theta_l d\chi_l \\ &= \prod_l \frac{-(1 + \Omega_l^{-1})}{2\pi} d\chi_l dj_l \\ &= \left[\prod_l \frac{-(1 + \Omega_l^{-1})}{2\pi} \right] d\Phi dJ dq dp \\ &\quad \times dQ_1 \cdots dQ_{L-2} dP_1 \cdots dP_{L-2} \end{aligned} \quad (\text{III.23})$$

where (Q_i, P_i) are the other canonical variables decoupled from the collective coordinates. From the third line to the forth line, we used that Jacobian equals one in canonical transformation. The part of the invariant measure which contributes to the excited states is

$$d\mu(\Phi, J; q, p) \propto d\Phi dJ dq dp = d\Phi dJ dE dt. \quad (\text{III.24})$$

Under EBK quantization condition (II.23), the k -th excited state becomes

$$\begin{aligned} |\tilde{\phi}_k\rangle &\propto \oint d\Phi \oint dt |\Phi, J; q, p\rangle e^{i\mathcal{T}(\Phi, J; q, p)} \\ &= \sum_{\{j_l\}} \oint d\Phi e^{i(J - \sum_l j_l)\Phi} \\ &\quad \times \oint dt e^{i\mathcal{T}_{\text{intr}}(t)} A_{j_l}(q, p) |\cdots; S_l, -S_l + j_l, \cdots\rangle \\ &\equiv \sum_{\{j_l\} \in (\sum j_l = J)} C_{j_l} |\cdots; S_l, -S_l + j_l, \cdots\rangle \end{aligned} \quad (\text{III.25})$$

where

$$C_{j_l} = \oint dt e^{i\mathcal{T}_{\text{intr}}(t)} A_{j_l}(q(t), p(t)). \quad (\text{III.26})$$

The definition of wave function in SPA can also be extended into ground state. The ground state is constructed by only energy minimum point in TDHFB phase space. In the energy minimum point, $|\Phi, J; q, p\rangle = |\text{HFB}\rangle$. Therefore, (III.25) becomes

$$|\tilde{\phi}_{g.s.}\rangle \propto \sum_{\{j_l\}} \oint d\Phi e^{i(J - \sum_l j_l)\Phi} |\text{HFB}\rangle, \quad (\text{III.27})$$

which is identical to the wave function of the particle number projection for HFB state.

IV. RESULTS

We apply ASCC+SPA to study the multi-level system in pairing model. The TDHFB degrees of freedom equals the number of single particle levels, including the constant of motion (pairing rotation). We consider various systems, two-level system, three-level system, and Pb isotope system in each subsection respectively. Next, we discuss the dynamics in non-integrable (more than three-level) system.

A. Confirmation in one-dimensional TDHFB system

The two-level system corresponds to one-dimensional TDHFB system after decoupling the global gauge angle Φ . We can obtain the obvious collective path from ASCC. The dynamics is exactly identical with adiabatic TDHFB (ATDHFB). We confirm that whether ASCC is identical to one-dimensional ATDHFB, and compare the difference with TDHFB, in numerical calculation.

In two-level system, the classical Hamiltonian in (III.8) is

$$\begin{aligned} \mathcal{H} &= \sum_{l=1,2} \epsilon_l \Omega_l (1 - \cos \theta_l) \\ &\quad - \frac{g}{4} \sum_{l=1,2} \Omega_l [\Omega_l (1 - \cos^2 \theta_l) + (1 - \cos \theta_l)^2] \\ &\quad - \frac{g}{2} \Omega_1 \Omega_2 \sqrt{(1 - \cos^2 \theta_{l_1})(1 - \cos^2 \theta_{l_2})} \cos(\chi_2 - \chi_1). \end{aligned} \quad (\text{IV.1})$$

If we define the canonical coordinate $\phi = \chi_2 - \chi_1$, it attributes to one-dimensional system. The conjugate momentum is $j = \frac{\partial \mathcal{L}}{\partial \phi} = \{\Omega_2(1 - \cos \theta_2) - \Omega_1(1 - \cos \theta_1)\}/4$. The ATDHFB indicates that the Hamiltonian can be expanded up to second order with respect to ϕ

$$\mathcal{H}(\phi, j) \approx V(j) + \frac{1}{2} B^{-1}(j) \phi^2, \quad (\text{IV.2})$$

where $V(j) = \mathcal{H}(\phi = 0, j)$ and $B^{-1}(j) = \left. \frac{\partial^2 \mathcal{H}}{\partial \phi^2} \right|_{\phi=0}$. We know that the BCS ground state corresponds to the potential minimum point with $\phi = 0$. If the pairing correlation is strong enough to bind the excited states in the collective potential, the adiabatic approximation is expected to be well because the states are localized in small ϕ region.

We study the system with equal degeneracy $\Omega_1 = \Omega_2 = 8$, $g/(\epsilon_2 - \epsilon_1) = 0.2$, and $N = 16$. In ASCC calculation, we set the mesh size of collective coordinate $dq = 0.01$. We compared the results of $|0_2^+\rangle$ and $|0_3^+\rangle$ in three different calculations. Fig. 2 and Fig. 3 correspond to the classical informations, which are classical trajectories in phase space and action integral \mathcal{T} as a function of time t . All of the mesh points obtained from ASCC are on the trajectories obtained from ATDHFB. For $|0_2^+\rangle$, the closed trajectory is well localized in the phase space and TDHFB calculation is almost the same as ASCC and ATDHFB calculations. For $|0_3^+\rangle$, the trajectory is still closed but near the transition point between closed trajectory and open trajectory. We can find ASCC and ATDHFB calculations have a small deviation from TDHFB calculation. The obtained excited 0^+ states from SPA are shown in Fig. 4. We show the occupation probability which is decomposed into $2n$ -particle- $2n$ -hole components. The results from ATDHFB and ASCC are identical within numerical error. Also, TDHFB calculation is very close to other two calculations.

Comparing with one-dimensional TDHFB calculation, we confirmed we succeed to describe the bound collective excited states by ASCC+SPA. For the bound states, whether the adiabatic approximation included or not hardly has influence. Even for the weakly bound states, the adiabatic approximation is reasonable approximation.

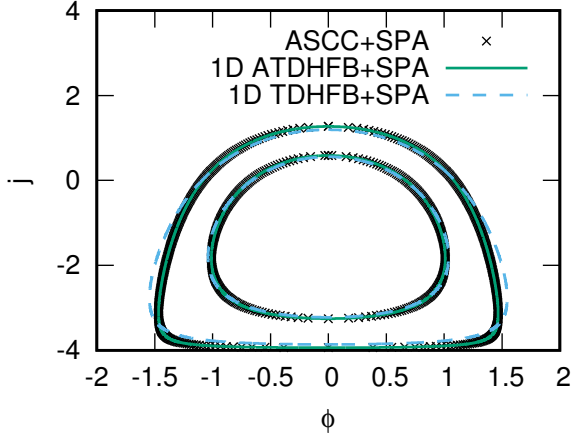


FIG. 2: Classical trajectories of $|0_2^+\rangle$ and $|0_3^+\rangle$ in two-dimensional phase space. The domain of phase space is $-\pi \leq \phi < \pi$, $-4 \leq j \leq 4$. Cross points, solid lines, and dashed lines correspond to the results from ASCC+SPA, one-dimensional ATDHFB+SPA, and one-dimensional TDHFB+SPA, respectively. In ASCC+SPA calculation, each cross point corresponds to each mesh point from ASCC calculation.

B. Three-level system

The simplest non-integrable system is three-level system. We have one trivial motion and two time-dependent degrees of freedom. We study the system with $\Omega_1 = \Omega_2 = \Omega_3 = 8$, $\epsilon_1 = -1$, $\epsilon_2 = 0$, $\epsilon_3 = 1.5$, and $g = 0.2$. The phase transition occurs at $g_c = 0.058$ when $N = 2\Omega_1 = 16$. We consider the even particle number chain from $N = 14$ to $N = 24$.

From moving-frame QRPA equation, we obtain three modes along the collective path (Fig. 5). Except zero mode, we choose the lowest mode as collective motion. Fig. 6 and Fig. 7 are occupation number in each single-particle level and collective potential in the collective path, respectively. We can find the occupation numbers become to integer at both end points. Furthermore, Hartree-Fock states always emerge at one of the end point.

Based on the information from ASCC calculation, we can obtain the excited states from SPA. Table I shows the excitation energies of first and second excited states in the collective subspace. Up to the second excitation, all of the energies are in their energy pockets shown in Fig. 7. Comparing the result from ASCC+SPA with exact solution, we can find all of the values are nicely reproduced. The slightly difference is that the values from ASCC+SPA are about 5% smaller than exact solution. Using the wave function of excited states, the pair additional transition strength is shown in Fig. 8. For intraband transition, comparing ASCC+SPA with exact solution, $B(P_{ad}; 0_1^+ \rightarrow 0_1^+)$ are almost identical and $B(P_{ad}; 0_2^+ \rightarrow 0_2^+)$ are about 10% ~ 20% small. For

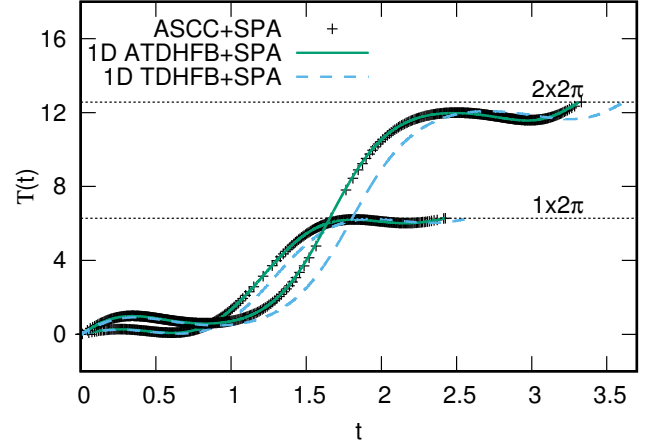


FIG. 3: Action integrals of $|0_2^+\rangle$ and $|0_3^+\rangle$ with respect to time t . Cross points, solid lines, and dashed lines correspond to the results from ASCC+SPA, one-dimensional ATDHFB+SPA, and one-dimensional TDHFB+SPA, respectively. Dotted lines are the values correspond to EBK quantization condition for $|0_2^+\rangle$ and $|0_3^+\rangle$. Based on Fig. 2, we calculated the action integrals on each trajectory from $(\phi, j) = (0, j_{\max})$ in clockwise direction.

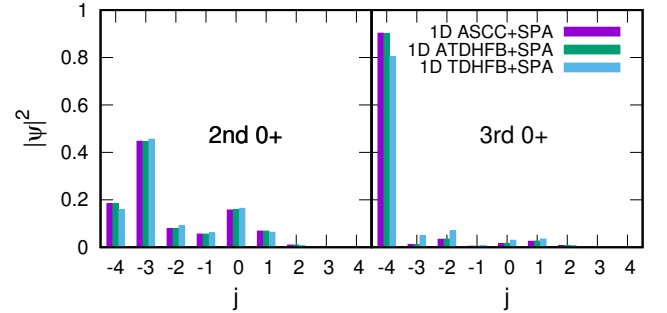


FIG. 4: Occupation probabilities of $|0_2^+\rangle$ and $|0_3^+\rangle$ as a function of j . The two vertical bars at each j from the left to the right represent the squared components of the wave functions from ASCC+SPA, ATDHFB+SPA and TDHFB+SPA calculations, respectively. The left end of the horizontal axis at $j = j_{\min}$ corresponds to a component with $(n_1, n_2) = (N, 0)$. The next at $j = j_{\min} + 1$ corresponds to the one with $(n_1, n_2) = (N - 2, 2)$, and so on.

interband transition, the strength from ASCC+SPA is much smaller than exact solution. It is difficult to reproduce the absolute value of exact solution. However, because the pairing correlation is enough strong in this case, the strength of intraband transition is dominant compared with interband transition. We can find that

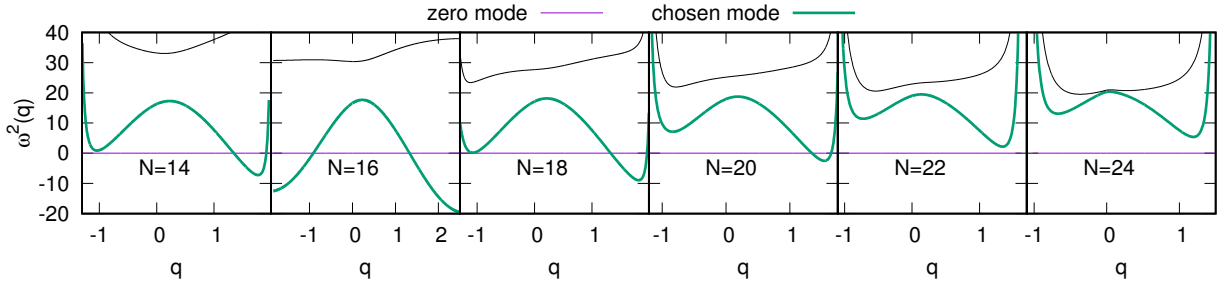


FIG. 5: Eigenvalues of moving-frame QRPA equation with respect to the collective coordinate q , from $N = 14$ to $N = 24$. Purple lines are spurious modes and green lines are chosen modes corresponding to the collective coordinate. In each panel, both edges correspond to the end points of the collective coordinate.

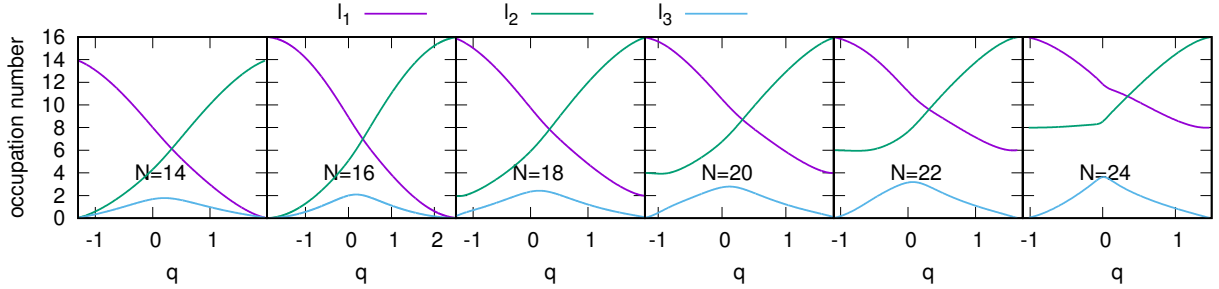


FIG. 6: Occupation numbers in each single-particle level with respect to the collective coordinate q , from $N = 14$ to $N = 24$. At the left end point of the collective coordinate in each panel, the configurations correspond to Hartree-Fock states.

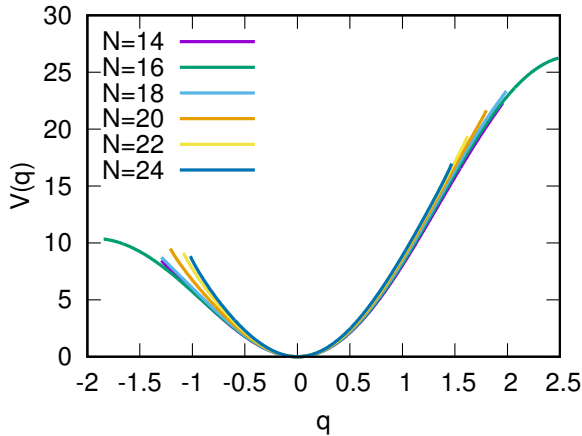


FIG. 7: Collective potential obtained from ASCC. We adjusted the energy minimum point as $V = 0$.

$B(P_{\text{ad}}; 0_1^+ \rightarrow 0_2^+)$ and $B(P_{\text{ad}}; 0_2^+ \rightarrow 0_1^+)$ are only about 1% of $B(P_{\text{ad}}; 0_1^+ \rightarrow 0_1^+)$ and $B(P_{\text{ad}}; 0_2^+ \rightarrow 0_2^+)$. Therefore, the interband transition can be regarded as reasonable. From all above results, we can conclude that ASCC+SPA reproduces exact solution quantitatively.

We need to discuss about the second excited state in

the collective path corresponds to not $|0_3^+\rangle$ and others, but $|0_4^+\rangle$. In pair additional transition $B(P_{\text{ad}}; k \rightarrow k')$ from exact solution, $B(P_{\text{ad}}; 0_2^+ \rightarrow 0_3^+)$ is 10 ~ 100 times smaller than $B(P_{\text{ad}}; 0_1^+ \rightarrow 0_3^+)$, and $B(P_{\text{ad}}; 0_1^+ \rightarrow 0_2^+)$ is in the same order with $B(P_{\text{ad}}; 0_1^+ \rightarrow 0_3^+)$. It indicates that $|0_3^+\rangle$ is also one-phonon state which belongs to the other collective path (black line in Fig. 5). About two-phonon states, we can easily find the candidates of two-phonon states from excitation energies and $B(P_{\text{ad}}; 0_1^+ \rightarrow 0_{\text{ex}}^+)$. In the candidates, $|0_4^+\rangle$ is the lowest two-phonon state and $B(P_{\text{ad}}; 0_2^+ \rightarrow 0_4^+)$ is much larger than $B(P_{\text{ad}}; 0_3^+ \rightarrow 0_4^+)$. Therefore, $|0_4^+\rangle$ is considered to be most appropriate corresponding to the second excited state in ASCC+SPA calculation.

C. Pb isotopes

With pairing model, we study neutron's pairing vibration in Pb isotope. We prepare the neutron's single-particle level between the magic number 82 and 126 (Table II). The coupling constant $g = 0.138(\text{MeV})$ is determined to reproduce the experimental pairing gap $\Delta(N) = \frac{(-1)^{N+1}}{2}(B(N+1) - 2B(N) + B(N-1))$ of ^{192}Pb . We consider the even-even system from ^{188}Pb to ^{194}Pb .

We have six TDHFB degrees of freedom in the system.

	N	14	16	18	20	22	24
0_2^+ one phonon	Exact	4.09	4.13	4.20	4.30	4.44	4.60
	ASCC+SPA	3.87	3.90	3.97	4.09	4.23	4.33
0_4^+ two phonon	Exact	7.65	7.71	7.88	8.15	8.49	8.74
	ASCC+SPA	7.42	7.42	7.60	7.92	8.26	8.47

TABLE I: Excitation energies of one-phonon and two-phonon states from exact solution and ASCC+SPA calculation. In a chosen collective subspace, one-phonon and two-phonon states correspond to the first and second excitation, respectively. In the whole Hilbert space, one-phonon and two-phonon states correspond to $|0_2^+\rangle$ and $|0_4^+\rangle$, respectively.

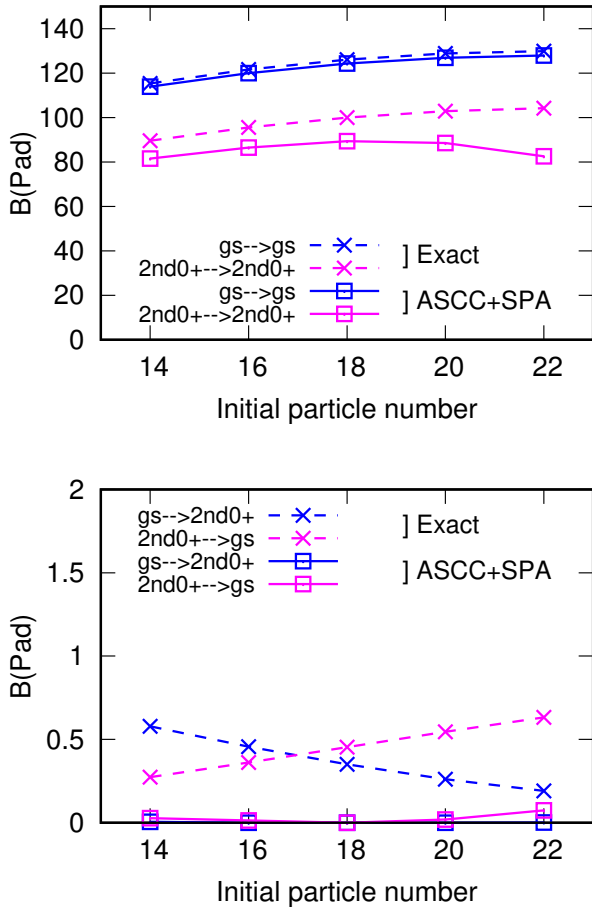


FIG. 8: The strength of pair additional transition $B(P_{ad}; k \rightarrow k') = |\langle N+2, k' | S^+ | N, k \rangle|^2$, from $N = 14$ to $N = 22$. Dashed lines are exact solution and solid lines are ASCC+SPA calculation. Horizontal line shows the particle number of the initial states. Upper panel shows the intraband transitions of $|0_1^+\rangle \rightarrow |0_1^+\rangle$ and $|0_2^+\rangle \rightarrow |0_2^+\rangle$, while lower panel shows the interband transition of $|0_1^+\rangle \rightarrow |0_2^+\rangle$ and $|0_2^+\rangle \rightarrow |0_1^+\rangle$.

orbit	$h_{9/2}$	$f_{7/2}$	$i_{13/2}$	$p_{3/2}$	$f_{5/2}$	$p_{1/2}$
energy(MeV)	-10.94	-10.69	-8.74	-8.44	-8.16	-7.45

TABLE II: Single-particle levels of Pb isotopes used in the calculation. The orbits are obtained from spherical Woods-Saxon potential with spin-orbit coupling.

Fig. 9 shows the eigenvalue of moving-frame QRPA equation. We can find there are one spurious mode and five vibrational modes. The collective modes we focus on did not cross with other modes in all panels, while the spacings of the collective modes and the second vibrational modes are small, especially for ^{194}Pb . Under the chosen collective modes, the occupation number is shown in Fig. 10. As with three-level system, the left end points correspond to HF state in all panels. The last neutrons are in $i_{13/2}$ from ^{188}Pb to ^{194}Pb . In the collective path, we can find $i_{13/2}$ and $p_{3/2}$ changes significantly, while other orbits don't change so much. The collective potential for each isotope is shown in Fig. 11. The depths of them are about 2.5 ~ 3.5(MeV) only enough to bind $|0_2^+\rangle$, one-phonon states. We can consider that the collectivity of pairing is small for excited states, even though their ground states are superfluid states.

We show the results of $|0_2^+\rangle$, one-phonon state in the collective coordinate, from SPA. Table II shows the excitation energy. The value is nothing to do with experimental value because pairing model is too simple to describe realistic nuclear interaction. We are only interested in the comparison between exact solution and ASCC+SPA. We can find the excitation energy from ASCC+SPA reproduce the exact solution as well as in three-level system. The pair additional transition strength is shown in Fig. 12. In this case, intraband transition is also dominant compared with interband transition. ASCC+SPA reproduces the tendency of exact solution and somewhat smaller than exact solution in all transition. However, the largest deviation, $B(P_{ad}; 1 \rightarrow 1)$, only has about 25% deficiency compared with exact solution. Therefore, ASCC+SPA is supposed to be available into complex realistic system.

In Pb isotope system, there are some problems remaining. Up to now, ASCC+SPA is only available when the excited states are bound in the collective potential. For example, we cannot calculate $|0_2^+\rangle$ in ^{186}Pb because the potential is too shallow to bound the excited state. Also, the whole collective path in ^{196}Pb cannot be obtained due to the complex eigenvalue of moving-frame QRPA equation in the collective path.

	^{186}Pb	^{188}Pb	^{190}Pb	^{192}Pb	^{194}Pb	^{196}Pb
Exact	2.58	2.44	2.34	2.25	2.20	2.15
ASCC+SPA	unbound	2.31	2.21	2.12	2.04	unbound

TABLE III: The same as Table. I but for Pb isotopes. The energies are given in units of MeV.

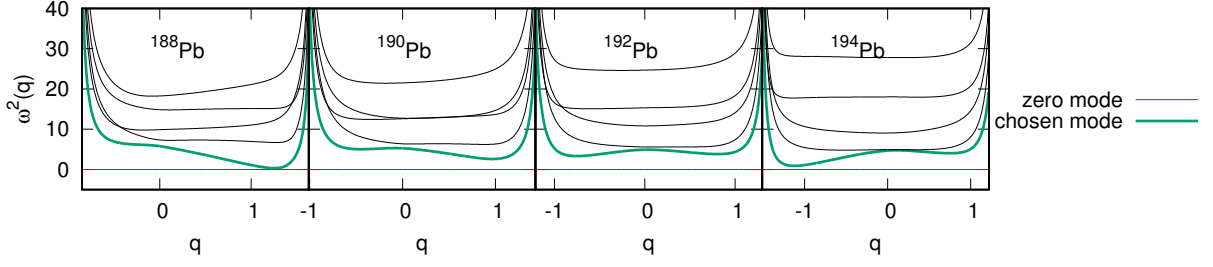


FIG. 9: The same as Fig. 5 but for Pb isotopes.

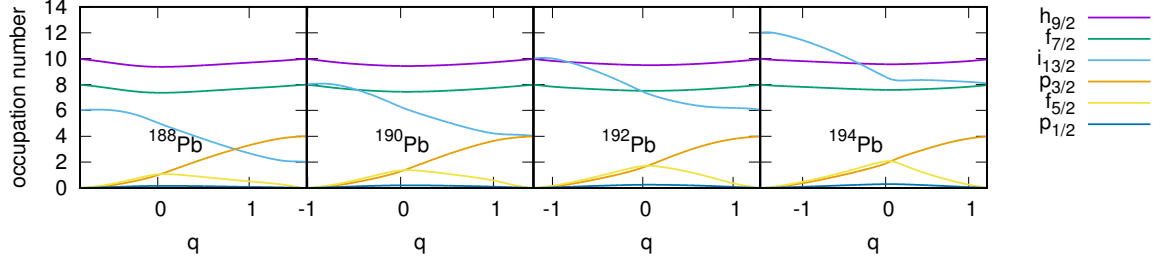


FIG. 10: The same as Fig. 6 but for Pb isotopes.

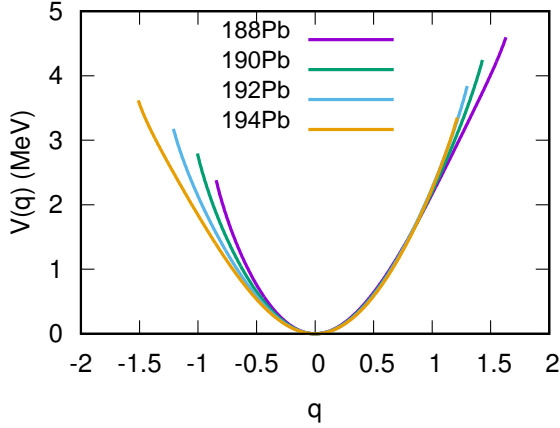


FIG. 11: The same as Fig. 7 but for Pb isotopes.

Finally, we discuss about whether the collective coordinate can be described by other physical quantities. The pairing gap Δ is the candidate of the collective coordinate, which is similar with $\alpha_{2\mu}$ in 5D Bohr Hamiltonian.

V. CONCLUSION

Based on the previous work which demonstrated the accuracy of SPA for the requantization of TDHFB dynamics, we combined ASCC with SPA to apply non-integrable system. We constructed the theoretical framework of ASCC+SPA for bound state, which is bound in the collective potential from ASCC. Besides the collective Hamiltonian, Q operator at each point of collective path is also necessary input information from ASCC.

We applied ASCC+SPA into multi-level pairing model. Because the global gauge angle Φ is a cyclic variable, the simplest non-integrable system is three level system. We investigated three level system and more realistic system, neutron's single particle levels from Pb isotope. In both cases, the low-lying excited 0^+ states reproduce exact solution well not only excitation energies but also wave functions. The pair transition calculation has no difficulty because we don't need to requantize the pair transition operators. This point overcomes the disadvantage in canonical requantization.

While the basic theoretical framework of ASCC+SPA was constructed, there are some problems remaining.

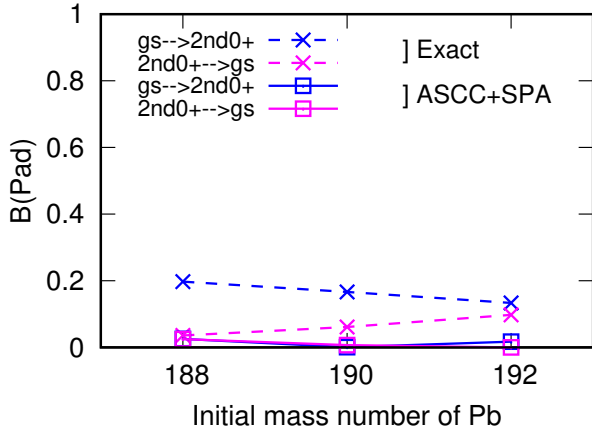
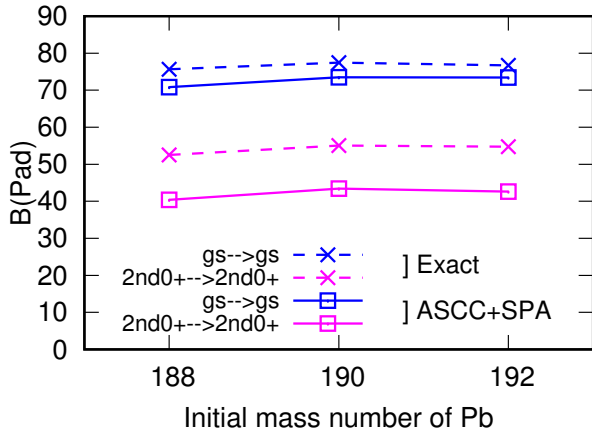


FIG. 12: The same as Fig. 8 but for Pb isotopes.

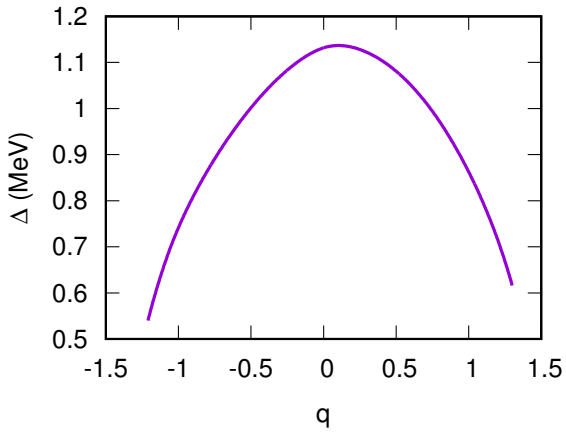


FIG. 13: Pairing gap as a function of collective coordinate in ^{192}Pb .