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Polaron dynamics with a multitude of Davydov D2 trial states
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Polaron dynamics with a multitude of Davydov D$_2$ trial states

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We propose an extension to the Davydov D$_2$ Ansatz in the dynamics study of the Holstein molecular crystal model with diagonal and off-diagonal exciton-phonon coupling using the Dirac-Frenkel time-dependent variational principle. The new trial state by the name of the “multi-D$_2$ Ansatz” is a linear combination of Davydov D$_2$ trial states, and its validity is carefully examined by quantifying how faithfully it follows the Schrödinger equation. Considerable improvements in accuracy have been demonstrated in comparison with the usual Davydov trial states, i.e., the single D$_1$ and D$_2$ Ansätze. With an increase in the number of the Davydov D$_2$ trial states in the multi-D$_2$ Ansatz, deviation from the exact Schrödinger dynamics is gradually diminished, leading to a numerically exact solution to the Schrödinger equation. © 2015 AIP Publishing LLC. [http://dx.doi.org/10.1063/1.4923009]

I. INTRODUCTION

Since the advent of the ultrafast laser spectroscopy, much attention has been devoted to the relaxation dynamics of photoexcited entities, such as polarons in inorganic liquids and solids, charge carriers in topological insulators, electron and hole trapping of semiconductor nanoparticles, and electron-hole pairs in light-harvesting complexes of photosynthetic organisms. In photosynthesis, it is suggested that strong dependence of electronic tunneling upon certain coordinated distortions of neighboring molecules in the formation of bound excited states. However, in the literature, little attention has been paid to the Hamiltonians containing the off-diagonal exciton-phonon coupling due to inherent difficulties to obtain reliable solutions, especially for the polaron dynamics. Early treatments of off-diagonal coupling include the Munn-Silbey theory, which is based upon a perturbative approach with added constraints on canonical transformation coefficients determined by a self-consistency equation. The global-local (GL) Ansätze formulated by Zhao et al. in the early 1990s, was later employed in combination with the dynamic coherent potential approximation (with the Hartree approximation) to arrive at a state-of-the-art ground-state wave function as well as higher eigenstates.

In the absence of an exact solution, various numerical approaches were developed in the past few decades, including the exact diagonalization (ED) and quantum Monte Carlo (QMC) simulation, variational method, and the method of relevant density matrix renormalization group (DMRG) and the variational exact diagonalization (VED), and the method of relevant coherent states. Most of these approaches were designed to probe the ground-state properties. For excited-state properties and dynamics of the polaronic systems, however, few of them provide a satisfactory resolution. For example, a time-dependent variant of DMRG, was developed to elucidate the polaron dynamics. Yet, it cannot accurately simulate the system dynamics from an arbitrary initial state, since high-lying excited states cannot be adequately described by DMRG. Fortunately, the variational approach is still effective in dealing with polaron dynamics so long as a proper trial wave function is chosen. Previously, static properties of the Holstein polaron have been examined using a series of trial wave functions based upon phonon coherent states, such as the Toyozawa Ansatz.
II. METHODOLOGY

The one-dimensional Holstein molecular crystal model for the exciton-phonon system can be described by the Hamiltonian below,

$$\hat{H} = \hat{H}_{\text{ex}} + \hat{H}_{\text{ph}} + \hat{H}_{\text{ex-ph}}^{\text{diag}} + \hat{H}_{\text{ex-ph}}^{\text{o.d.}},$$

where $\hat{H}_{\text{ex}}$, $\hat{H}_{\text{ph}}$, and $\hat{H}_{\text{ex-ph}}^{\text{o.d.}}$ correspond to the exciton Hamiltonian, bath (phonon) Hamiltonian, and exciton-phonon coupling Hamiltonian defined as

$$\hat{H}_{\text{ex}} = -J \sum_{n} \hat{a}_{n}^\dagger (\hat{a}_{n+1} + \hat{a}_{n-1}),$$
$$\hat{H}_{\text{ph}} = \sum_{q} \omega_{q} \hat{b}_{q}^\dagger \hat{b}_{q},$$
$$\hat{H}_{\text{ex-ph}}^{\text{diag}} = -g \sum_{n,q} \omega_{q} \hat{a}_{n}^\dagger \hat{a}_{n} (e^{i\varphi_{n}} \hat{b}_{q} + e^{-i\varphi_{n}} \hat{b}_{q}^\dagger),$$
$$\hat{H}_{\text{ex-ph}}^{\text{o.d.}} = \frac{1}{2} \phi \sum_{n,q} \omega_{q} \left[ \hat{a}_{n}^\dagger \hat{a}_{n+1} [e^{i\omega_{n}} (e^{i\varphi_{n}} - 1) \hat{b}_{q} + \text{H.c.}] + \hat{a}_{n}^\dagger \hat{a}_{n-1} [e^{i\omega_{n}} (1 - e^{-i\varphi_{n}}) \hat{b}_{q} + \text{H.c.}] \right],$$

where H.c. denotes the Hermitian conjugate, $\omega_{q}$ is the phonon frequency at the momentum $q$, $\hat{a}_{n}$ ($\hat{a}_{n}^\dagger$) is the exciton creation (annihilation) operator for the $n$th molecule, and $\hat{b}_{q}$ ($\hat{b}_{q}^\dagger$) is the creation (annihilation) operator of a phonon with the momentum $q$.

The parameters $J$, $g$, and $\phi$ represent the transfer integral, diagonal coupling strength, and off-diagonal coupling strength, respectively, and $N = 16$ is the number of sites in the molecular ring of the Holstein polaron. In this paper, a linear dispersion phonon band is assumed,

$$\omega_{q} = \omega_{0} \left[ 1 + W(2|q|/\pi) - 1 \right],$$

where $\omega_{0}$ denotes the central energy of the phonon band, $W$ is the band width between 0 and 1, and the momentum is set to be $q = 2\pi l/N$ with $l = \frac{l}{2} + 1, \ldots, N - \frac{1}{2}$.

A trial state, termed as the “Davydov multi-D$_{2}$ Ansatz,” is adopted

$$|D_{2}^{M}(t)\rangle = \sum_{n=1}^{N} \hat{a}_{n}^\dagger |0\rangle_{\text{ex}} \sum_{i=1}^{M} \psi_{i,n}(t) \times \exp \left[ \sum_{q} \left( \lambda_{i,q} \hat{b}_{q}^\dagger - \lambda_{i,q}^* \hat{b}_{q} \right) \right] |0\rangle_{\text{ph}},$$

where $\hat{a}_{n}^\dagger$ ($\hat{a}_{n}$) is the creation (annihilation) operator of an exciton at the $n$th site, $\hat{b}_{q}^\dagger$ ($\hat{b}_{q}$) is the creation (annihilation) operator of a phonon with momentum $q$, and the variational parameters $\psi_{i,n}$ and $\lambda_{i,q}$ denote the exciton probability and phonon displacement, respectively. Moreover, $n$ and $i$ represent the site number in the molecular ring and the index of the coherent superposition state, respectively.

The equations of the motion are derived for the variational parameters $\psi_{i,n}$ and $\lambda_{i,q}$ by adopting the Dirac-Frenkel variational method, in which the Lagrangian $L$ is formulated as

$$L = \langle D_{2}^{M}(t) | \frac{i\hbar}{2} \frac{\partial}{\partial t} - \hat{H} | D_{2}^{M}(t) \rangle$$
$$= \frac{i\hbar}{2} \left[ \langle D_{2}^{M}(t) | \frac{\partial}{\partial t} | D_{2}^{M}(t) \rangle - \langle D_{2}^{M}(t) | \frac{\partial}{\partial t} | D_{2}^{M}(t) \rangle \right]$$
$$- \langle D_{2}^{M}(t) | \hat{H} | D_{2}^{M}(t) \rangle.$$
From this Lagrangian, the equations of the motion for α and its time derivative \( \dot{\alpha}(t) \) can be obtained,

\[
\frac{d}{dt} \left( \frac{\partial L}{\partial \dot{\alpha}^*} \right) - \frac{\partial L}{\partial \alpha^*} = 0,
\]

where \( \alpha \) is one of the variational parameters \( \psi_{i,n} \) and \( \lambda_{i,q} \) in Eq. (5). Details on derivation of the equations of the motion for the Holstein polaron dynamics with a multi-D2 Ansatz are given in Appendix A.

As shown in Fig. 2, the initial states of the Holstein polaron are prepared for the diagonal coupling case with the coupling strength \( g > \phi = 0 \) in (a) and the off-diagonal coupling case with the coupling strength \( \phi > g = 0 \) in (b). In order to avoid singularity, a little noise uniformly distributed from \([ - \varepsilon, \varepsilon]\) is added with \( \varepsilon = 10^{-5} \) in the initial state for both \( \psi_{i,n} \) and \( \lambda_{i,q} \). For each set of the coefficients \( W, g, J, \) and \( \phi \) defined in Eqs. (2) and (4), more than 100 initial states are used in the simulations. A power-law relation between the Central Processing Unit (CPU) time and the multiplicity \( M \) has been found in our time-dependent variational approach, and an exponent of 2.8 indicates that the time complexity of the program is \( O(N^3) \). Even for the largest value of the multiplicity in our paper, i.e., \( M = 64 \), the memory usage of the multi-D2 Ansatz is 256 MB (million bytes), still bearable for the computation. Though both the CPU time and memory usage of the multi-D2 Ansatz are much larger than those of the single D2 Ansatz (less than 1 hour for the CPU time and 0.1 MB for the memory usage), the multi-D2 Ansatz has substantially improved the accuracy on variational dynamics. The energy of the system \( E_{\text{total}} = E_{\text{ex}} + E_{\text{ph}} + E_{\text{diag}} + E_{\text{off}} \) is calculated based on the multi-D2 Ansatz in Eq. (5).

\[
E_{\text{ex}} = \langle D^M_2 | \hat{H}_{\text{ex}} | D^M_2 \rangle
\]

\[
= -J \sum_{i,j} \sum_{n} \psi_{i,n} (\phi_{i,n+1} + \phi_{i,n-1}) S_{j,i},
\]

\[
E_{\text{ph}} = \langle D^M_2 | \hat{H}_{\text{ph}} | D^M_2 \rangle
\]

\[
= \sum_{i,j} \sum_{n} \psi_{i,n}^* \psi_{i,n} \sum_{q} \omega_q \lambda_{j,q} \lambda_{i,q} S_{j,i},
\]

\[
E_{\text{diag}} = \langle D^M_2 | \hat{H}_{\text{diag}} | D^M_2 \rangle = -g \sum_{i,j} \sum_{n} \psi_{i,n}^* \psi_{i,n} \sum_{q} \omega_q \left( e^{iqn} \lambda_{i,q} + e^{-iqn} \lambda_{j,q} \right) S_{j,i},
\]

\[
E_{\text{off}} = \langle D^M_2 | \hat{H}_{\text{off}} | D^M_2 \rangle = \frac{1}{2} \sum_{i,j} \sum_{n} \omega_q S_{j,i}
\]

\[
\times \left( \left| \psi_{i,n} \psi_{i,n+1} \right| e^{iqn} (e^{iq} - 1) \lambda_{i,q} + \text{H.c.} \right) \psi_{j,n}^* \psi_{j,n+1} \left( (e^{iq} - 1) \lambda_{j,q} + \text{H.c.} \right)
\]

where \( S_{j,i} = \langle \lambda_{i,j} | \lambda_{j,i} \rangle \) is the Debye-Waller factor defined as

\[
S_{j,i} = \exp \left( -\frac{1}{2} \left( |\lambda_{i,j}|^2 + |\lambda_{j,i}|^2 \right) \right).
\]

The normalization of the wave function \( \text{Norm} = \langle D^M_2 | D^M_2 \rangle \) is also calculated to test its conservation.

Furthermore, the exciton probability \( P_{\text{ex}}(t,n) \) and the phonon displacement \( X_{\text{ph}}(t,n) \) are also calculated for the dynamics of the Holstein polaron. First, the reduced single-exciton density matrix \( \rho_{mn}(t) = \text{Tr}[\rho(t) \hat{a}_m^\dagger \hat{a}_n] \) is obtained by solving the coupled equations of variational parameters, where \( \rho(t) = |D^M_2\rangle \langle D^M_2| \) is the full density matrix at zero temperature. After substituting the trial state \( |D^M_2(t)\rangle \) of Eq. (5), the reduced density matrix is then derived as

\[
\rho_{mn}(t) = \sum_{i,j} \psi_{i,m}^* \psi_{i,n} S_{j,i}.
\]

Thus, the exciton probabilities \( P_{\text{ex}}(t,n) = \rho_{mn}(t)(n = 1, 2, \ldots, N) \) are obtained from the diagonal elements of the reduced density matrix. The phonon displacement \( X_{\text{ph}}(t,n') \) in real space at the \( n' \)th site is calculated by

\[
X_{\text{ph}}(t,n') = \langle D^M_2 | b_{n'}(t) + b_{n'}^\dagger(t) | D^M_2 \rangle
\]

\[
= \frac{1}{\sqrt{N}} \sum_q e^{iqn'} \left[ \sum_{i,j} \sum_n \lambda_{i,q} \psi_{i,n}\psi_{i,n} S_{j,i} \right]
\]

\[
+ \frac{1}{\sqrt{N}} \sum_q e^{-iqn'} \left[ \sum_{i,j} \sum_n \lambda_{j,q} \psi_{j,n}^* \psi_{i,n} S_{j,i} \right].
\]

Optical spectroscopy is another important aspect of the polaron dynamics, as it can provide valuable information on various correlation functions. In this work, the linear absorption spectra for the polaron dynamics calculated with different Ansätze will be studied to check the validity of these trial wave functions. The linear absorption spectra \( F(\omega) \) can be obtained by the Fourier transformation,

\[
F(\omega) = \frac{1}{\pi} \text{Re} \int_0^\infty F(t) e^{i\omega t} dt,
\]
where \( F(t) \) is the autocorrelation function of the exciton-phonon system, defined as

\[
F(t) = \langle \hat{\rho}_e(0) | e^{-iHt} \hat{\rho}_e e^{-iHt} \hat{\rho}_e | 0 \rangle_{\text{ph}}
\]

\[
= \langle \hat{\rho}_e(0) | e^{-iHt} \hat{\rho}_e | 0 \rangle_{\text{ph}}
\]

with the polarization operator

\[
\hat{\rho} = \mu \sum_n (\hat{a}_n^\dagger(0) | 0 \rangle_{\text{ex}} | 0 \rangle_{\text{ex}} + | 0 \rangle_{\text{ex}} | 0 \rangle_{\text{ex}} \hat{a}_n^\dagger(0)).
\]

Details on how to calculate the linear absorption spectra of a one-dimensional exciton-phonon system from the multi-D, single D, and single D\(_1\) Ansätze are given in Appendix C.

Finally, the validity of our Ansatz, Eq. (5), will be closely scrutinized. Assuming the trial wave function \( | \Psi(t) \rangle = | \Psi(t) \rangle \) at the moment \( t \), a deviation vector \( | \delta(t) \rangle \) is then introduced to quantify the accuracy of the variational method,

\[
| \delta(t) \rangle = \frac{\partial}{\partial t} | \Psi(t) \rangle - \frac{\partial}{\partial t} | \Psi(t) \rangle.
\]

Thus, deviation from the exact Schrödinger dynamics can be indicated by the amplitude of the deviation vector \( \Delta(t) = | \delta(t) \rangle | \delta(t) \rangle \). In order to view the deviation in the parameter space \( (W, J, g, \phi) \), a dimensionless relative deviation \( \sigma \) is calculated as

\[
\sigma = \frac{\max \{ \Delta(t) \}}{\text{mean} \{ E_{\text{ph}}(t) \}}, \quad t \in [0, t_{\text{max}}],
\]

where the phonon energy \( E_{\text{ph}}(t) \) is the main energy of the system and is almost the same as the amplitude of the time derivative of the wave function,

\[
N_{\text{ex}}(t) = \sqrt{\frac{-\langle \frac{\partial}{\partial t} | \Psi(t) \rangle, \frac{\partial}{\partial t} | \Psi(t) \rangle \rangle}{\langle \hat{H}^2 | \hat{\rho}_e \hat{\rho}_e | 0 \rangle_{\text{ex}}}}
\]

\[
\approx \Delta E,
\]

since \( \langle E \rangle = \langle \hat{D}^2(t) \hat{H} | \hat{\rho}_e \hat{\rho}_e | 0 \rangle_{\text{ex}} \approx 0 \) in this paper.

### III. NUMERICAL RESULTS

#### A. Diagonal coupling

The long-time behavior of the Holstein polaron dynamics is described by Eq. (7). Fig. 3 shows the evolution of the system energies, including the total energy \( E_{\text{total}} \), the phonon energy \( E_{\text{ph}} \), the exciton energy \( E_{\text{ex}} \), and the exciton-phonon interaction energy \( E_{\text{diag}} \), for the diagonal coupling case with \( J = 0.1, W = 0.5, \) and \( g = 1 \). For \( N = 16 \) sites in the molecular ring, the Ansatz is formed from a superposition of \( M = 16 \) D\(_2\) wave functions, and the initial state shown in Fig. 2(a) is used. The periodicity of the system energies is given by

\[
T = 4\omega_0/\pi, \quad \text{in perfect agreement with the expectation of } N/4W.
\]

The finding that \( E_{\text{ph}} \approx -E_{\text{diag}} \) and \( E_{\text{ex}} \approx E_{\text{total}} \approx 0 \) shows that the total energy is conserved in the Dirac-Frenkel variational dynamics.

The exciton probability \( P_{\text{ex}}(t, n) \) and the phonon displacement \( X_{\text{ph}}(t, n) \) from the multi-D\(_2\) Ansatz are compared with those from the single D\(_2\) and D\(_1\) Ansätze. The latter can be written as

\[
| \Delta(t) \rangle = \sum_{n=1}^{N} \phi_n(t) | a_n^\dagger | 0 \rangle_{\text{ex}}
\]

\[
\times \exp \left[ \sum_q \left( \lambda_{n,q} \beta_q^\dagger - \lambda_{n,q}^* \beta_q \right) \right] | 0 \rangle_{\text{ph}}.
\]

where the displacement coefficient \( \lambda_{n,q} \) is not only dependent on the moment \( q \) but also on the site index \( n \) in the molecular ring. As referred in the "Introduction," while the D\(_1\) Ansatz is effective in the diagonal coupling case, it fails to describe the polaron dynamics of the Holstein model with off-diagonal coupling.

In Fig. 4, the time evolution of the exciton probability \( P_{\text{ex}}(t, n) \) and the phonon displacement \( X_{\text{ph}}(t, n) \) are shown in a weak-coupling case with \( g = 0.1 \) calculated by the D\(_2\) Ansatz [panels (a) and (b)], the D\(_2\)-16 Ansatz [panels (c) and (d)], and the D\(_1\) Ansatz [panels (e) and (f)]. Similarly, time-dependent behaviors of both \( P_{\text{ex}}(t, n) \) and \( X_{\text{ph}}(t, n) \) are found to be almost the same in the multi-D\(_2\) and the single D\(_1\) Ansätze, but at variance with those in the single D\(_2\) Ansatz. Moreover, the propagation of the exciton wave packets can be found in Fig. 4(c) with a velocity \( v \approx \omega_0/2\pi \), consistent with that obtained by the D\(_1\) Ansatz in Fig. 4(e).
The behavior of the Holstein polaron in the intermediate diagonal coupling regime is shown in Fig. 5 at the diagonal coupling strength $g = 0.2$. Similar time-dependent pattern in $P_{ex}(t,n)$ and $X_{ph}(t,n)$ is spotted in the multi-D$_2$ Ansatz and the D$_1$ Ansatz, but not in the single D$_2$ Ansatz. The speed of the localized exciton wave packets $v \approx \omega_0/4\pi$ is then measured from both Figs. 5(c) and 5(e) to be nearly half of that in the weak coupling case of $g = 0.1$. It suggests that the velocity $v$ is inversely proportional to the diagonal coupling strength $g$. Moreover, similar phonon propagation patterns are found in (d) and (f), including the sound waves and the movement induced by the exciton-phonon interaction. The latter is found to be with the same velocity as that of the exciton wave packets.

The results of the variational dynamics in the strong coupling case with $g = 0.4$ are shown in Figs. 6(a)–6(f), corresponding to the D$_2$, D$_2^{M=16}$, and D$_1$ Ansätze, respectively. Different from the weak and intermediate coupling cases in Figs. 4 and 5, the exciton probabilities $P_{ex}(t,n)$ and phonon displacements $X_{ph}(t,n)$ obtained by these three kinds of the trial wave functions are nearly identical, indicating that the D$_2$, multi-D$_2$, and D$_1$ Ansätze are effective in the strong diagonal coupling case.

Using the amplitude of the deviation vector $\Delta(t)$, validity of the multi-D$_2$ Ansatz defined in Eq. (5) is investigated for various values of the multiplicity $M$, as shown in Fig. 7. The amplitude $\Delta(t)$ is almost constant, and the time-averaged value $\langle \Delta(t) \rangle$ monotonically decreases with $M$. It indicates that the multi-D$_2$ trial state approaches the exact solution to the Schrödinger equation when $M$ is sufficiently large. For comparison, $\Delta(t)$ obtained by the D$_1$ Ansatz is also plotted, the time average of which is smaller than that obtained from the single D$_2$ Ansatz ($M = 1$), consistent with our previous contention that D$_1$ Ansatz is more accurate than D$_2$ Ansatz in the diagonal coupling regime. Interestingly, $\langle \Delta(t) \rangle$ calculated from the multi-D$_2$ Ansatz with $M = 16$ is 0.05, much smaller than 0.14 from the D$_1$ Ansatz and 0.40 from the single D$_1$ Ansatz, showing the superiority of the multi-D$_2$ Ansatz.

Among the three trial states, the validity test of the multi-D$_2$ states is comprehensively performed for various values of the diagonal coupling strength $g$ and transfer integral $J$ in the diagonal coupling only regime. In Fig. 8, the relative deviation $\sigma$ defined in Eq. (17) is displayed as a function of the diagonal coupling strength $g$ for various values of $M$ for the case of $J = 0.1, W = 0.5$, and $\phi = 0$. As $M$ increases, the relative error $\sigma$ decreases, especially for the weak coupling case of $g = 0.1$. For comparison, $\sigma$ calculated from the D$_1$ Ansatz is also shown, with $\sigma$ obviously larger than that of the multi-D$_2$ Ansatz with $M = 32$, further supporting the superiority of the multi-D$_2$ Ansatz over the D$_1$ state. Moreover, $\sigma \approx 0.01$ at $M = 32$ indicates that the variational method based on the multi-D$_2$ Ansatz is possible to be numerically exact in the limit of $M \to \infty$, where “numerically exact” means the relative error $\sigma = 0$ within numerical errors.
FIG. 5. Time evolution of the exciton probability \( P_{\text{ex}}(t, n) \) and the phonon displacement \( X_{\text{ph}}(t, n) \) obtained by the \( D_2 \), \( D_{M=16} \), and \( D_1 \) trial states are displayed in (a)-(f) for \( J = 0.1, W = 0.5, g = 0.2, \) and \( \phi = 0 \).

The behavior of the relative error \( \sigma \) as a function of the transfer integral \( J \) is also investigated for several values of \( M \). As shown in Fig. 9, \( \sigma \) monotonically increases with \( J \). When \( J \) becomes much larger than the reorganization energy, the wave function approaches a plane wave, which is difficult to be described by superpositions of coherent states. With an increase in \( M \), however, the relative deviation \( \sigma \) is monotonically reduced indicating the improved efficiency of the multi-\( D_2 \) Ansatz even in the case with a large transfer integral. Moreover, the results show that the multi-\( D_2 \) Ansatz is at least as accurate as the \( D_1 \) Ansatz if \( M = 32 \) superpositions are used.

B. Off-diagonal coupling

In this section, we probe the dynamics of the Holstein polaron in the off-diagonal coupling regime via the multi-\( D_2 \) Ansatz, in comparison with those obtained by the single \( D_2 \) and the single \( D_1 \) Ansätze. The initial state is given the form shown in Fig. 2(b). For simplicity, only the off-diagonal coupling is considered in the simulations. Time evolution of the exciton probability \( P_{\text{ex}}(t, n) \) and phonon displacement \( X_{\text{ph}}(t, n) \) obtained by the single \( D_1 \), the single \( D_2 \), and the \( D_{M=16} \) Ansätze are displayed. In Figs. 10(a) and 10(b) for the \( D_1 \) Ansatz, one can find that self-trapping occurs at the 8th and 9th sites, at variance with the expected delocalization of the Holstein dynamics in the off-diagonal coupling case. It is consistent with the previous impression that \( D_1 \) Ansatz fails to describe the dynamics of the Holstein polaron in the off-diagonal coupling regime.

In contrast, the spread of the exciton is found in the results obtained by the single \( D_2 \) and multi-\( D_2 \) Ansätze. However, as shown in Figs. 10(c) and 10(d), the wave function obtained by the single \( D_2 \) Ansatz is still localized, different from those obtained by the multi-\( D_{M=16} \) Ansatz shown in the Figs. 10(e) and 10(f). It indicates that the multi-\( D_2 \) Ansatz is more effective than the single \( D_2 \) Ansatz for depicting the Holstein polaron dynamics in the off-diagonal coupling case. Moreover, one can find three different stages of the exciton motion in Figure 10(e), separated by the characteristic time \( t_1 \approx 2\pi/\omega_0 \) and \( t_2 \approx 8\pi/\omega_0 \). When \( t < t_1 \), the exciton is self-trapped in the initial state, pointing to a localized exciton state. After that, the exciton starts to spread over the molecular ring with the velocity \( v \approx \omega_0/\pi \). Until \( t \geq t_2 \), a uniform distribution of the exciton wave packets emerges, indicating that the exciton is in a delocalized state.

Via the relative deviation \( \sigma \) defined in Eq. (17), the validity of the multi-\( D_2 \) Ansatz can be further confirmed. In Fig. 11, the relative deviation \( \sigma \) is plotted as a function of \( 1/M \) for the off-diagonal coupling only case with \( \phi = 1.0 \). Both the
For the strong diagonal coupling strength $g = 0.4$, time evolution of the exciton probability $P_{ex}(t, n)$ and the phonon displacement $X_{ph}(t, n)$ obtained by the $D_2$, $D_{16}^M$, and $D_1$ Ansätze are displayed in (a)-(f) at $J = 0.1$, $W = 0.5$, and $\phi = 0$.

The amplitude of the deviation vector $\Delta(t)$ from the Schrödinger equation is plotted as a function of the time $t$ with the unit $2\pi/\omega_0$ at $J = 0.1$, $W = 0.5$, $g = 1.0$, and $\phi = 0$. Lines represent the results obtained from the multi-$D_2$ Ansatz with various values of $M$ and the stars denote those from the $D_1$ Ansatz.

The relative deviation $\sigma$ defined in Eq. (17) is displayed as a function of the diagonal coupling strength $g$ at $J = 0.1$, $W = 0.5$, and $\phi = 0$. The lines with dots represent the results obtained from the multi-$D_2$ Ansatz with various values of $M$ and the dashed line denotes those from the $D_1$ Ansatz. The size of the molecular ring is $N = 16$. 

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FIG. 9. The relative deviation $\sigma$ in a 16-site molecular ring is displayed as a function of the transfer integral $J$ in the case of $g = 1, W = 0.5, \phi = 0$. The lines with dots represent the results obtained from the multi-D$_2$ Ansatz with various values of $M$ and the dashed line denotes those from the D$_1$ Ansatz.

As $M$ increases, the relative deviation $\sigma$ decreases and approaches zero as $M$ goes to infinity. For example, the value of $\sigma(M = 60) = 0.26$ is much smaller than $\sigma(M = 1) = 0.67$. According to the fitting in the inset, the relationship $\sigma \sim M^{-\nu}$ is revealed with the exponent $\nu = 0.29(1)$, further confirming the prediction $\sigma = 0$ in the limit of $M \to \infty$. Hence, it can be concluded that the variational method based on the multi-D$_2$ Ansatz can be numerically exact ($\sigma = 0$) in both the diagonal and off-diagonal coupling regimes. Since any quantum state of a system of multiple oscillators (boson modes) can be represented by a continuous superposition of coherent states (often referred to as the unit decomposition property of the aforementioned states), there is a good chance that the multi-D$_2$ Ansatz is numerically exact, i.e., it provides exact results in the $M \to \infty$ limit. However, it remains unclear how practical is the above statement, since the values of the multiplicity $M$, needed for the Ansatz to converge to the exact solution of the dynamical Schrödinger equation might be unrealistically large. The above question will be addressed elsewhere.

Finally, accuracy of the multi-D$_2$ Ansatz is quantified for the parameter regime $0 \leq g, \phi \leq 1$, in comparison with that of the single-D$_2$ Ansatz, as shown in Fig. 12. The influence of the excitonic initial state is also taken into account. Figs. 12(a) transfer integral $J$ and the phonon bandwidth $W$ are set to be zero. As $M$ increases, the relative deviation $\sigma$ decreases and approaches zero as $M$ goes to infinity. For example, the value of $\sigma(M = 60) = 0.26$ is much smaller than $\sigma(M = 1) = 0.67$. According to the fitting in the inset, the relationship $\sigma \sim M^{-\nu}$ is revealed with the exponent $\nu = 0.29(1)$, further confirming the prediction $\sigma = 0$ in the limit of $M \to \infty$. Hence, it can be concluded that the variational method based on the multi-D$_2$ Ansatz can be numerically exact ($\sigma = 0$) in both the diagonal and off-diagonal coupling regimes. Since any quantum state of a system of multiple oscillators (boson modes) can be represented by a continuous superposition of coherent states (often referred to as the unit decomposition property of the aforementioned states), there is a good chance that the multi-D$_2$ Ansatz is numerically exact, i.e., it provides exact results in the $M \to \infty$ limit. However, it remains unclear how practical is the above statement, since the values of the multiplicity $M$, needed for the Ansatz to converge to the exact solution of the dynamical Schrödinger equation might be unrealistically large. The above question will be addressed elsewhere.

Finally, accuracy of the multi-D$_2$ Ansatz is quantified for the parameter regime $0 \leq g, \phi \leq 1$, in comparison with that of the single-D$_2$ Ansatz, as shown in Fig. 12. The influence of the excitonic initial state is also taken into account. Figs. 12(a)
FIG. 11. The relative deviation $\sigma$ from the multi-D$_2$ Ansatz is displayed as a function of $1/M$ for the off-diagonal coupling case with the strength $\phi = 1$, and other parameters $J = W = g = 0$ are set. The size of the molecular ring is $N = 16$. In the inset, the relationship $\sigma \sim M^{-\nu}$ is displayed on a log-log scale and the dashed line represents a power-law fit.

and 12(b) correspond to the one-site occupied initial state shown in Fig. 2(a) and Figs. 12(c) and 12(d) correspond to the two-site occupied initial states shown in Fig. 2(b). For each initial state, two different values of the transfer integral $J = 0.1$ and 0.5 are used. Our results show that the D$_{M=16}$ Ansatz deviates little from the exact solutions of the time-dependent Schrödinger equation in the whole parameter regime for both the two initial states and the cases with small and large transfer integrals, thereby further confirming the validity of the variational method. Moreover, the significant improvement of the validity for the multi-D$_2$ Ansatz from the single D$_2$ Ansatz is found especially in the regime with weak diagonal and off-diagonal coupling, lending support to the high accuracy of the Ansatz.

C. Absorption spectra

Besides the relative deviation, the validity of the Ansätze in providing reliable dynamical information can also be gauged by the accuracy of optical spectra, as analytical expressions for the absorption and fluorescence spectra are well-known if the transfer integral $J$ and the phonon bandwidth $W$ are negligible. In this study, the set of the parameters $J = 0.1, W = 0.1, g = 0.4$, and $\phi = 0$ is used, and the Huang-Rhys factor $S$ is then calculated according to the relaxation energy defined by

$$E_r \equiv \int_{-\infty}^{\infty} \frac{C_{00}(\omega)}{\omega} d\omega = \sum_q g_q^2 \omega_q \equiv S\omega_0, \quad (20)$$

where $\omega_0$ is the central energy of the phonon band, $g_q = g = 0.4$ is the diagonal coupling, and $\omega_q$ is the frequency at the moment $q$. From this equation, we can obtain $S = 2.56$ corresponding to the diagonal coupling strength $g = 0.4$. Two types of initial states, one-site-occupied excited state and the excitation with a Gaussian-type distribution spanned on 7 sites, have been applied in the study of optical spectra. To facilitate comparisons, spectral maxima are normalized to unity.

FIG. 12. The relative deviation $\sigma$ is displayed for the single D$_2$ and multi-D$_2$ Ansätze as a function of the diagonal coupling strength $g$ and off-diagonal coupling strength $\phi$. In (a) and (b), the exciton at $t = 0$ is created on a single site $n = N/2$, while in (c) and (d), a two-site occupied state is used as the initial state.
Linear absorption spectra of a 16-site one-dimensional ring of a coupled exciton-phonon system are displayed in (a) for the single D$_2$ Ansatz and in (b) for the D$_M$-16 Ansatz. Two kinds of the initial states including the one-site occupied and Gaussian occupied exciton distributions are used in the simulations, and the Huang-Rhys factor is $S = 2.56$. For comparison, the numerical results obtained by the single D$_1$ Ansatz and the fitting of a Poisson distribution are given in (b) with the solid circles and bars, respectively. A rescaled factor is applied to normalize the spectral maxima for facilitating comparison.

Three trial states including the single D$_1$, the single D$_2$, and the multi-D$_2$ Ansätze are investigated, as they differ in terms of the variational parameters in describing the phonon behavior. Linear absorption spectrum is a very useful indicator of the Ansatz validity in the investigation of the dynamics of a polaron system. For the single D$_2$ Ansatz, the phonon displacement is only described by one set of variational parameters, leading to the lack of exciton-phonon correlation between exciton and phonon, and eventually to the absence of long-range correlation in the autocorrelation function and an inaccurate description of optical spectra. Only for the cases with the one-site occupied initial state under strong diagonal coupling and small $J$ where the exciton is localized, as shown by the black solid line in Fig. 13(a), the inability of the single D$_1$ Ansatz is alleviated. As for the red line, where the initial electronic excitation adopts a Gaussian distribution, the spectrum even exhibits negative values around $\omega = -3.5\omega_0$. In contrast, the single D$_1$ and the multi-D$_2$ Ansätze guarantee the long-range exciton-phonon correlation, therefore can provide accurate absorption spectra. Fig. 13(b) shows similarly accurate spectra for both single D$_1$ and multi-D$_2$ Ansätze.

According to the Huang-Rhys theory, the phonon sidebands at zero temperature follow a Poisson distribution:

$$F(\omega) = \exp(-S) \sum_{n=0}^{\infty} \frac{S^n}{n!} \delta (\omega + E_n - n\omega). \quad (21)$$

From Eq. (21), the left most sideband, $n = 0$, corresponding to the zero-phonon line, should be located at $\omega = -S\omega_0$, i.e., $\omega = -2.56\omega_0$ consistent with our result $\omega = -2.64\omega_0$ as shown in Fig. 13(b). What is more, the tallest peaks at $n = 1$ and $2$ show almost same height, in agreement with the prediction that the tallest of phonon sidebands should be at $n = S - 1 = 1.56$ when $S \gg 1$. Further, using a fitting method, the bar plot of the Poisson distribution with the parameter $\lambda = 2.2$ is given, consistent with our spectra obtained from time-dependent evolution of single D$_1$ and multi-D$_2$ Ansätze. The slight deviation of the Poisson parameter $\lambda$ from the Huang-Rhys factor $S = 2.56$ is due to the nonzero values of $W$ and $J$.

**IV. CONCLUSION**

Using the multi-D$_2$ Ansatz as the trial state, we have systematically studied the dynamics of a one-dimensional Holstein polaron with simultaneous diagonal and off-diagonal exciton-phonon coupling via the Dirac-Frenkel time-dependent variational approach. Compared to the single D$_2$ Ansatz, the multi-D$_2$ counterpart is much more sophisticated and contains more flexible variational parameters, leading to superior quality simulations of polaron dynamics. Special attention is paid to testing the validity of our time-dependent variational approach by quantifying how closely the trial state follows the Schrödinger equation. Linear absorption spectra derived from the trial state are also studied as a sensitive indicator of the Ansatz validity in the investigation of polaron dynamics.

Our numerical results show considerable improvements in accuracy of polaron dynamics by the multi-D$_2$ Ansatz, in comparison with the usual single D$_1$ and D$_2$ trial states. In the diagonal coupling regime, the multi-D$_2$ Ansatz is found to be at least as accurate as the single D$_1$ Ansatz for various values of the diagonal-coupling strength $g$ and the transfer integral $J$ and remarkably better than the single D$_2$ Ansatz. In the off-diagonal coupling regime, however, the multi-D$_2$ Ansatz is shown to be much more potent in depicting the Holstein polaron dynamics than the single D$_2$ Ansatz, while the single D$_1$ Ansatz fails completely. As the number of superposed states $M$ increases, one can find visible decays of the relative deviation $\sigma$ in the weak diagonal coupling regime as well as the
off-diagonal coupling regime, confirming respectable accuracies of the multi-D$_2$ Ansatz. Moreover, $\sigma = 0$ is predicted by the numerical fitting in the limit of $M \to \infty$, inferring that the Dirac-Frenkel time-dependent variational approach based on the multi-D$_1$ is possible to be numerically exact.

The single Davydov D$_1$ Ansatz is a trial state with sufficient flexibilities to handle accurately quantum dynamics from the celebrated spin-boson model to large light-harvesting complexes in photosynthesis. Very recently, a systematic coherent-state expansion of the ground state wave function that is based on the Davydov D$_1$ Ansatz, which we shall call the “multi-D$_1$ Ansatz,” is developed for a number of models. It is a generalization of a variational wave function originally proposed by Silbey and Harris and also an extension of the hierarchy of translationally invariant Ansätze proposed by Zhou et al. The results of the quantum phase transition obtained from the multi-D$_1$ Ansatz are more accurate than those from the single D$_1$ Ansatz, and they are in agreement with DMRG and ED results, showing the superiority of the multi-D$_1$ Ansatz. The successful application of the multi-D$_2$ Ansatz in the Holstein polaron dynamics points to the possibility that the multi-D$_1$ Ansatz is not only valid for studying static and also an extension of the multi-D$_2$ Ansatz has convincingly shown that even a relatively simple wave function such as the Davydov D$_2$ Ansatz, when used in an expandable superposition, can still produce superior results. If the D$_2$ Ansatz were to be replaced by the more sophisticated D$_1$ trial state and applied in a multitude as demonstrated here, much better results can be expected on simulating quantum dynamics of many-body systems. Work in this direction is now in progress.

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APPENDIX A: TIME EVOLUTION OF THE MULTI-D$_2$ TRIAL STATE

As mentioned in Sec. II “Methodology,” the time evolution for the multi-D$_2$ Ansatz can be derived by employing Dirac-Frenkel time-dependent variational method. According to the definition of the multi-D$_2$ Ansatz in Eq. (5) and the Dirac-Frenkel variational principle, the variational parameters $\psi_{i,n}(t)$ and $\lambda_{i,q}(t)$ should obey

$$\frac{d}{dt} \left( \frac{\partial L}{\partial \psi_{i,n}} \right) - \frac{\partial L}{\partial \psi_{i,n}} = 0,$$

$$\frac{d}{dt} \left( \frac{\partial L}{\partial \lambda_{i,q}} \right) - \frac{\partial L}{\partial \lambda_{i,q}} = 0,$$

(A1)

where $L$ is the Lagrangian defined in Eq. (6). Inside the Lagrangian, the first term reads as following:

$$\langle D^M_2(t) | \frac{\partial}{\partial t} | D^M_2(t) \rangle = E_{ex} + E_{ph} + E_{diag} + E_{off},$$

(A3)

where $E_{ex}$, $E_{ph}$, $E_{diag}$, and $E_{off}$ denote the energies of the exciton, phonon, diagonal coupling term, and off-diagonal coupling term, respectively, which can be calculated based on Eq. (8).

Time partial derivatives of $\lambda_{i,q}(t)$ and $\psi_{i,n}(t)$ are then obtained by substituting Eqs. (A2) and (A3) into Eq. (A1), which are

$$-i \sum_{i} \frac{\partial}{\partial q} \psi_{i,n} \Rightarrow \frac{1}{2} \sum_{i,j} \left( \frac{\partial \lambda_{i,j}^*}{\partial q} \lambda_{i,j} + \frac{\partial \lambda_{i,j}}{\partial q} \lambda_{i,j}^* \right) S_{i,j}$$

$$= \sum_{i} \psi_{i,n+1} + \psi_{i,n-1} \Rightarrow \sum_{i} \psi_{i,n} \sum_{q} \omega_{q} \lambda_{k} \lambda_{i} \Rightarrow \sum_{i} \omega_{q} \left( e^{iqn} \lambda_{i} + e^{-iqn} \lambda_{i}^* \right) S_{k,i}$$

$$- \frac{1}{2} \sum_{i} \sum_{q} \omega_{q} \psi_{i,n} \left( e^{iqn} (e^{i\lambda_{i,q} - 1}) + e^{-iqn} (e^{-i\lambda_{i,q} - 1}) \right) S_{k,i}$$

$$- \frac{1}{2} \sum_{i} \sum_{q} \omega_{q} \psi_{i,n} \left( e^{iqn} (1 - e^{-i\lambda_{i,q}}) + e^{-iqn} (1 - e^{i\lambda_{i,q}}) \right) S_{k,i}$$

(A4)
where $S_{k,i}$ is the Debye-Waller factor defined in Eq. (9). Unfortunately, both Eqs. (A4) and (A5) contain the coupled partial time derivatives of $\lambda_{i,q}(t)$ and $\psi_{i,q}(t)$ with the form $\sum_i A_i \psi_{n,i} + \sum_{i,q} A_2 \lambda_{i,q} + \sum_{i,q} A_3 \lambda_{i,q}^* = B$, where $A_1, A_2, A_3,$ and $B$ are coefficient vectors, unrelated to the partial time derivatives of the variational parameters. By numerically solving these linear equations at each time $t$, one can calculate the values of $\psi_{i,q}$ and $\lambda_{i,q}$ accurately. The 4th order Runge-Kutta method is then adopted for the time evolution of the Holstein polaron, including the energies $E_{\text{total}}(t) = E_{\text{ex}} + E_{\text{ph}} + E_{\text{diss}} + E_{\text{off}}$, the normalization $\text{Norm}(t)$, the exciton probability $P_{\text{ex}}(t,n)$, and phonon displacement $X_{\text{ph}}(t,n)$.

Finally, the amplitude of the deviation vector $\tilde{\Delta}(t)$ defined in Eq. (15) is calculated as

$$\Delta^2(t) = \langle \tilde{\Delta}(t) \rangle$$

$$= \sum_n \sum_{i,j} \left[ \sum_q \left( A_{jq}^* B_{jq} + B_{jq}^* A_{jq} \right) \right] S_{ji}$$

$$\times \left\{ \sum_q \left( A_{iq}^* B_{iq} + B_{iq}^* A_{iq} \right) \right\}$$

where the matrices $A_{jq}$ and $B_{jq}$ are, respectively, defined as

$$A_{jq} = \frac{1}{N} \psi_{j,n}(t)$$

$$- i \frac{1}{2} \psi_{j,n}(t) \left[ \lambda_{j,q}(t) A_{jq}^* + A_{jq} \lambda_{j,q}^*(t) \right]$$

$$+ J \frac{1}{N} \psi_{j,n+1}(t) + \psi_{j,n-1}(t) + g \phi \psi_{j,n}(t) \omega_q e^{i\omega_n \lambda_{j,q}}$$

and

$$B_{jq} = i \phi \psi_{j,n}(t) \lambda_{j,q}^*(t) - \psi_{j,n}(t) \omega_q \lambda_{j,q} + g \phi \psi_{j,n}(t) \omega_q e^{-i\omega_n \lambda_{j,q}}$$

$$- i \frac{1}{2} \phi \psi_{j,n+1}(t) \omega_q e^{-i\omega_n \lambda_{j,q}}$$

$$- i \frac{1}{2} \phi \psi_{j,n-1}(t) \omega_q e^{-i\omega_n \lambda_{j,q}}.$$

**APPENDIX B: ENERGY CONSERVATION**

In the diagonal coupling case of $J = 0.1, W = 0.5$, and $g = 1$, the total energy $E_{\text{total}}(t)$ and the normalization of the wave function $\text{Norm}(t)$ are plotted as a function of the time $t$ in Fig. 14 for the precision test of the multi-D2 Ansatz with $M = 16$. One can find that the deviations of $E_{\text{total}}$ and $\text{Norm}$ from the initial values are negligibly small, suggesting that the numerical results obtained by the Dirac-Frenkel variational dynamics based on the multi-D2 trial states are reliable.

Besides, the dynamic behavior of the Holstein polaron for the off-diagonal coupling case is also investigated at $J = 0, W = 0, g = 0$, and $\phi = 1$. As shown in Fig. 15, aperiodic behaviors of the system energies are found, consistent with the prediction of the long period time $T \to \infty$ due to the vanishing of the band width $W = 0$. $E_{\text{total}}(t) = E_{\text{ph}} + E_{\text{off}} \approx 0$ shows the system total energy is conserved. In the inset, the normalization $\text{Norm}(t)$ is also displayed for the conservativeness test.

**APPENDIX C: LINEAR ABSORPTION**

Combining Eqs. (13) and (14), the autocorrelation function is derived

$$F(t) = \mu^2 \sum_n \sum_m \text{ph} \langle 0 | \text{ex} | \langle 0 | \hat{\alpha}_m e^{-i\hat{H}t} \hat{\alpha}_n^* \rangle_{\text{ex}} \rangle_{\text{ph}}.$$  

(C1)

Using the periodic condition of the Hamiltonian $\hat{H}$, one has

$$\sum_m \left( \sum_n \text{ph} \langle 0 | \text{ex} | \langle 0 | \hat{\alpha}_m e^{-i\hat{H}t} \hat{\alpha}_n^* \rangle_{\text{ex}} \rangle_{\text{ph}} \right)$$

$$= \sum_m \left( \sum_n \text{ph} \langle 0 | \text{ex} | \langle 0 | \hat{\alpha}_m e^{-i\hat{H}t} \hat{\alpha}_n^* \rangle_{\text{ex}} \rangle_{\text{ph}} \right)$$

(C2)

Substituting Eq. (C2) into Eq. (C1), one can obtain

$$F(t) = \mu^2 N \sum_m \left( \sum_n \text{ph} \langle 0 | \text{ex} | \langle 0 | \hat{\alpha}_m e^{-i\hat{H}t} \hat{\alpha}_n^* \rangle_{\text{ex}} \rangle_{\text{ph}} \right),$$

(C3)

where $e^{-i\hat{H}t} \hat{\alpha}_n^* \langle 0 | \text{ex} \rangle_{\text{ph}}$ is the time evolution of wave function from the initial state $\hat{\alpha}_n^* \langle 0 | \text{ex} \rangle_{\text{ph}}$, which can be approximated...
In the diagonal coupling case, the total energy of the system $E_{\text{total}}$ and the normalization of the wave function Norm are plotted as a function of the time $t$ for a molecular ring with $N = 16$ sites. The time unit $2\pi/\omega_0$ denotes the vibrational period of the phonon. The parameters including the transfer integral $J = 0.1$, phonon energy bandwidth $W = 0.5$, diagonal coupling strength $g = 1$, and off-diagonal coupling strength $\phi = 0$ are set.

For the single $D_2$ trial state, the time evolution of wave function from the initial state can be approximated to

$$e^{-i\hat{H}_t} |\psi_0(0)\rangle = |\psi_0(t)\rangle.$$

The autocorrelation $F(t)$ of the multi-$D_2$ Ansatz is then calculated by substituting Eq. (C4) into Eq. (C3),

$$F(t) = \mu^2 \sum_{ij} \sum_n \sum_m \psi_{i,m}(t)_{\text{ph}} \langle 0 | x 0_{\text{ph}} | 0 \rangle.$$

For the single $D_1$ trial state, the time evolution of wave function is then calculated by substituting Eq. (C6) into Eq. (C5),

$$e^{-i\hat{H}_t} |\psi_0(0)\rangle = |\psi_0(t)\rangle.$$

Finally, the single $D_1$ trial state is used to calculate the time evolution of the wave function

$$e^{-i\hat{H}_t} |\psi_0(0)\rangle,$$

and the autocorrelation $F(t)$ is then obtained by

$$F(t) = \mu^2 \sum_{ij} \sum_n \sum_m \psi_{i,m}(t)_{\text{ph}} \langle 0 | x 0_{\text{ph}} | 0 \rangle.$$

by a Davydykov trial state, for example, by the Multi-$D_2$ trial state,

$$e^{-i\hat{H}_t} |\psi_0(0)\rangle = |\psi_0(t)\rangle.$$

The total energy of the system $E_{\text{total}}$ and the normalization of the wave function Norm are plotted as a function of the time $t$ for a molecular ring with $N = 16$ sites. The time unit $2\pi/\omega_0$ denotes the vibrational period of the phonon. The parameters including the transfer integral $J = 0.1$, phonon energy bandwidth $W = 0.5$, diagonal coupling strength $g = 1$, and off-diagonal coupling strength $\phi = 0$ are set.

For the single $D_2$ trial state, the time evolution of wave function from the initial state can be approximated to

$$e^{-i\hat{H}_t} |\psi_0(0)\rangle = |\psi_0(t)\rangle.$$

The autocorrelation $F(t)$ of the multi-$D_2$ Ansatz is then calculated by substituting Eq. (C4) into Eq. (C3),

$$F(t) = \mu^2 \sum_{ij} \sum_n \sum_m \psi_{i,m}(t)_{\text{ph}} \langle 0 | x 0_{\text{ph}} | 0 \rangle.$$

For the single $D_1$ trial state, the time evolution of wave function is then calculated by substituting Eq. (C6) into Eq. (C5),

$$e^{-i\hat{H}_t} |\psi_0(0)\rangle = |\psi_0(t)\rangle.$$


