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<p>Highlights</p>	<p>◇量子位相推定アルゴリズムによる量子化学計算では、求めたい電子状態の真の波動関数にできるだけ近い近似波動関数を事前に準備することが計算効率を上げる鍵となる。</p> <p>◇断熱量子計算アルゴリズムの一種である断熱状態生成法（ASP 法）を用い、求めたい電子状態の真の波動関数を効率的に生成するための実用的な計算条件を明らかにした。</p> <p>◇今後、量子化学の分野において ASP 法と量子位相推定を組み合わせる研究が広く展開されることが期待される。</p>
<p>Description</p>	<p>【概要】</p> <p>研究チームは、量子コンピュータ上で原子・分子の波動関数の精度を向上させることができる断熱状態生成法（adiabatic state preparation, ASP 法）の実用的な計算条件を、初めて明らかにしました。</p> <p>ASP 法は、複雑な電子構造を持つ分子の量子化学計算を量子コンピュータ上で効率的に行うための有力手法の 1 つであると考えられていましたが、具体的な計算条件がほとんど調べられておらず、実用的な手段とは言えない状況でした。ASP 法の実用的な計算条件を初めて明らかにした本研究成果は、量子コンピュータによる量子化学計算を実際の化学研究に役立てるための大きな一歩と言えます。</p> <p><研究の背景></p> <p>近年、暗号に利用する大きな桁数の素因数分解など、スパコンなどの古典コンピュータでは「問題サイズ」に対して指数関数的に計算時間が増えてしまう特定の課題に対して、多項式時間内の計算を可能にする量子コンピュータの研究が活発に行われています。これらの課題のなかでも、原子・分子のシュレーディンガー方程式※3 を近似的に解き、電子状態を明らかにする精密な量子化学計算は、量子コンピュータの近い将来の計算ターゲットとして注目されています。</p> <p>量子化学計算の主な手法として、全配置間相互作用法（full configuration interaction; full-CI 法）と呼ばれる計算法があります。古典コンピュータで full-CI 法を採用した場合、分子サイズに対して計算コストが指数関数的に増大します。一方、量子コンピュータでは量子位相推定という量子アルゴリズムを用いることにより、計算コストが分子サイズに対して多項式時間内の増大で収まるようになることが理論的に示されています。量子位相推定を用いた full-CI 計算では「射影測定」と呼ばれる測定を行い、インプットとして用いた近似波動関数を真の波動関数に射影する方法によって求めたい電子状態の波動関数とエネルギーを計算します。量子位相推定でどの電子状態の波動関数が得られるかは近似波動関数と真の波動関数がどれだけ似ているかに依存します。もしも事前に準備した近似波動関数が真の波動関数から大きく異なっていれば、求めたい電子状態のエネルギーを得るまで同じ計算を何回も実行する必要があり、量子コンピュータによる計算高速化の利点が損なわれてしまいます。そのため、近似波動関数の精度を上げる手法開発は、量子位相推定に基づく量子化学計算の実用化に必須な研究課題と言えます。</p> <p>これまで、量子コンピュータ上に生成した近似波動関数の精度を上げる手法はさまざま</p>

まなものが研究されていますが、本研究グループは ASP 法に注目しました。ASP 法は固有関数が簡単に準備できるシンプルなハミルトニアンから真のハミルトニアンへと徐々に変化させていくことにより真の波動関数を得る手法です。ASP 法はハミルトニアンを変化させるスピードを十分に遅くすることにより良い近似波動関数が得られる、求めたい電子状態に関する詳細な事前知識を必要としないなどの大きな利点があります。しかし、どれくらいの時間をかけてハミルトニアンを変化させればよいか、ハミルトニアンの時間変化をどのような関数形で表現すればよいか、ASP 法のスタート時の波動関数としてどのようなものを用いればよいかなど、これまで ASP 法の具体的な計算条件がほとんど調べられておらず、実用的な手段とは言えない状況でした。

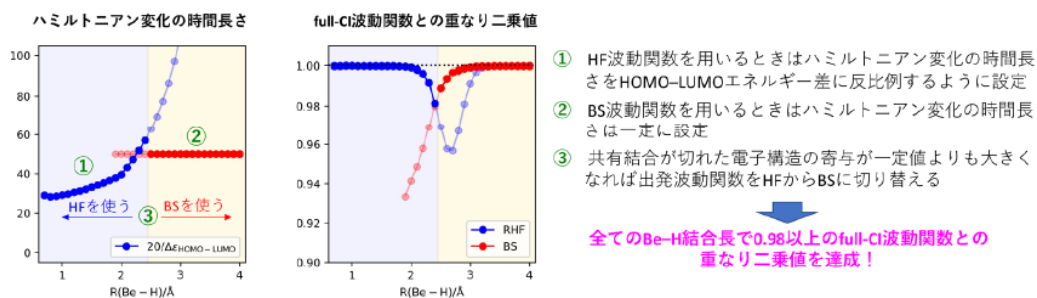
<研究の内容>

同研究グループは ASP 法の実用的な計算条件を探るために、①ハミルトニアンの時間変化を表現する関数として、従来用いられてきた一次関数だけでなく、三角関数などの非線形関数を試す、②ハミルトニアンを変化させる時間の長さを、分子軌道エネルギーから推定する、③ASP 法の初期波動関数として、従来用いられてきた Hartree-Fock (HF) 波動関数だけでなく、broken-symmetry (BS) 波動関数と呼ばれる波動関数を用いる、など、ASP 法に汎用性を持たせるために、計算条件をさまざまに変えて数値シミュレーションを行いました。

その結果、以下の 4 点が明らかになりました。

- (I) HF 波動関数からスタートしたとき、ハミルトニアンの時間変化はサイン関数を用いて表現することで最も良い近似波動関数を得られる。
- (II) HF 波動関数からスタートしたときはハミルトニアンを変化させる時間の長さは HOMO-LUMO エネルギー差に反比例するように設定すればよい。
- (III) 共有結合が切れた構造では BS 波動関数を用いることでより少ないステップ数で良い波動関数を生成できる。
- (IV) BS 波動関数を用いた場合、ハミルトニアンを変化させる時間の長さは分子構造によらず一定でよい。

さらに、初期波動関数として HF 波動関数と BS 波動関数のどちらを用いればよいかを簡単に判定する手法も提案しました (図)。これらの研究成果により、ASP 法の実用的計算条件が初めて明らかになりました。



図：BeH₂分子の2つのBe-H結合を同時に伸長した構造でのASP法の数値シミュレーション結果。左図はASP法でハミルトニアンを変化させる時間の長さを、右図はfull-CI波動関数との重なり二乗値（重なり二乗値は1に近ければ近いほど波動関数の近似精度が高いことを表す）を示す。図の横軸はBe-H結合長を表す。左図中の領域①、②、及び境界③に対する設定の説明を、右端に示す。

<期待される効果・今後の展開>

精密な量子化学計算は創薬や材料開発など、さまざまな関連分野でも非常に重要な役割を担っています。これまで報告されてきた量子位相推定によるfull-CI計算に関する理論的研究のほとんどは、良い近似波動関数が簡単に得られることを前提条件としていましたが、この前提条件をいかにして満足させるかは自明ではありません。本研究により、ASP法を用いて良い近似波動関数を生成するための実用的な道筋が初めて示されました。今後、精密な量子化学の分野においてASP法と量子位相推定を組み合わせる研究が広く展開され、量子コンピュータを用いた理論計算に基づく新たな材料設計などへ応用されることが期待されます。

‘量子コンピュータ上での量子化学計算の効率向上へ！分子の波動関数を生成するASP法の実用化に大きな一歩’ 大阪公立大学.

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Adiabatic state preparation of correlated wave functions with nonlinear scheduling functions and broken-symmetry wave functions

Kenji Sugisaki ^{1,2,3✉}, Kazuo Toyota¹, Kazunobu Sato ^{1✉}, Daisuke Shiomi ¹ & Takeji Takui ^{1,4✉}

Adiabatic state preparation (ASP) can generate the correlated wave function by simulating the time evolution of wave function under the time-dependent Hamiltonian that interpolates the Fock operator and the full electronic Hamiltonian. However, ASP is inherently unsuitable for studying strongly correlated systems, and furthermore practical computational conditions for ASP are unknown. In quest for the suitable computational conditions for practical applications of ASP, we performed numerical simulations of ASP in the potential energy curves of N_2 , BeH_2 , and in the C_{2v} quasi-reaction pathway of the Be atom insertion to the H_2 molecule, examining the effect of nonlinear scheduling functions and the ASP with broken-symmetry wave functions with the S^2 operator as the penalty term, contributing to practical applications of quantum computing to quantum chemistry. Eventually, computational guidelines to generate the correlated wave functions having the square overlap with the complete-active space self-consistent field wave function close to unity are discussed.

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Quantum computing and quantum information processing (QC/QIP) is one of the most innovative research fields in the current central science¹ and it has a potential to bring a paradigm shift in chemistry research. Among the diverse topics in the field of QC/QIP, sophisticated quantum chemical calculations have attracted attention as the near-future applications of quantum computers. Quantum chemical calculations are based on the Schrödinger equation that is a fundamental equation in quantum mechanics, and methods for accurate quantum chemical calculations potentially pave the way toward predictive quantum chemistry. Variationally best possible wave functions within the Hilbert space spanned by the basis set being used can be obtained by employing the full-configuration interaction (full-CI) calculations. However, the computational cost of full-CI methods scales exponentially against the number of basis functions relevant to the system size under study, and it easily reaches astronomical figures even for small molecules^{2,3}.

To date, two major approaches for the quantum chemical calculations on quantum computers have been widely investigated, namely quantum phase estimation (QPE)-based full-CI calculations and a variational quantum eigensolver (VQE). The QPE-based approach was proposed in 2005⁴. It is based on the quantum simulations of the time evolution of an approximated wave function and projective measurement to the eigenstate of a given Hamiltonian. The quantum circuit for the QPE-based full-CI is too deep to be executed on noisy intermediate-scale quantum (NISQ) devices currently available. The QPE-based approach, however, is expected to be a powerful tool when fault-tolerant quantum computing (FTQC) becomes available, because the computational cost of the QPE-based full-CI scales polynomially against the system size, and therefore exponential improvement of the computational cost scaling from the classical counterpart is guaranteed theoretically. In the QPE-based methods the time evolution of wave functions should be simulated conditionally on the ancillary qubit, but this requirement is recently removed by the appearance of the approach based on the Bayesian phase difference estimation (BPDE) algorithm⁵. VQE is a quantum–classical hybrid algorithm, and it utilizes parametrized quantum circuits to prepare correlated wave functions and computes energy expectation values by statistical sampling of the measurement outcome⁶. Classical computers are used to perform the variational optimizations of parameters relevant to the quantum circuits. VQE has been extensively studied from both the experimental and theoretical sides, because the parametrized quantum circuit used for wave function preparation is usually very shallow enough to be executed on NISQ devices. It should be noted that, however, the computational cost scaling of the VQE-based approaches has not been solely elucidated yet, and it is unclear whether quantum chemical calculations can be accelerated by using VQE. For example, VQE calculations with the unitary coupled cluster with singles and doubles (UCCSD) ansatz scale polynomial. The approximate UCCSD calculations, however, can also be done with the polynomial cost on a classical computer, by solving the amplitude equation of the similarity transformed Hamiltonian. Thus, the computational cost scaling becomes polynomial vs. polynomial for the UCCSD calculations on quantum and classical computers. Also, because the full-CI wave function contains an exponentially large number of variables, solving the full-CI using VQE with naïve implementation scales exponential. Recent reviews on quantum chemical calculations on quantum computers including QPE and VQE can be found elsewhere^{7–13}.

In the QPE-based full-CI calculations, the preparation of a “good” approximated wave function having sufficiently large overlap with the exact wave function of the target electronic state is crucial. This is because the probability of which eigenenergy of

the electronic state can be obtained in the QPE is proportional to the square overlap between the approximated and the exact wave functions. If the approximated wave function has exponentially small overlap with the eigenfunction of the target electronic state, an exponentially large number of QPE experiments are required to acquire the correct results, which spoils the advantages of a quantum speedup. In the equilibrium geometry of typical closed-shell singlet molecules and open-shell high-spin molecules carrying no unpaired electrons of spin- β , the Hartree–Fock (HF) wave function $|\Psi_{\text{HF}}\rangle$ dominantly contributes to the full-CI wave function of the electronic ground state, and the use of $|\Psi_{\text{HF}}\rangle$ is generally a good choice. In the low-spin states of open-shell molecules, by contrast, the wave function is approximated by the linear combination of several Slater determinants so as to be an eigenfunction of the S^2 operator, and the overlap between $|\Psi_{\text{HF}}\rangle$ and $|\Psi_{\text{full-CI}}\rangle$ becomes small. We demonstrated that the symmetry-adapted configuration state function (CSF) can have a large overlap with the full-CI wave function of open-shell low-spin systems, and proposed quantum circuits to prepare the $|\Psi_{\text{CSF}}\rangle$ on a quantum computer^{14,15}. We also reported an approach to generate multi-configurational wave functions on quantum computers without performing any post-HF calculations¹⁶, which is suitable for the study of the systems with intermediate open-shell characters like molecules under covalent bond dissociations. These approaches can effectively consider static (nondynamical) electron correlation effects.

Although these approaches are useful to treat electronic states of open-shell molecules, the overlap with the full-CI wave function becomes small when dynamical electron correlation effects are also significant. Molecules having electronic structures too complicated to deal with facile approaches like density functional theory (DFT) are naturally one of the main targets of sophisticated quantum chemical calculations, and the development of theoretical methods to generate correlated wave functions considering both static and dynamical electron correlation effects on quantum computers is an important task. Note that one of the anticipated usage of VQE is the preparation of approximate wave functions used as the input for QPE¹⁷, but recent numerical simulations of VQE of the C_{2v} quasi-reaction pathway of $\text{Be} + \text{H}_2 \rightarrow \text{BeH}_2$ reaction revealed that the variational parameter optimization converges very slowly for strongly correlated systems¹⁸.

Noticeably, recent years have witnessed that these promising approaches are relevant to innovative development in simulating quantum systems, such as Hamiltonian simulation. Adiabatic state preparation (ASP)⁴ is an approach to generate correlated wave functions based on an adiabatic theorem¹⁹. ASP belongs to an adiabatic quantum algorithm^{20,21}, in which the wave function of the ground state of a problem Hamiltonian H_P is generated adiabatically, by starting from the ground state wave function of an initial Hamiltonian H_I and slowly varying the Hamiltonian by using a scheduling function $s(t)$ in Eq. 1 from 0 to 1.

$$H(t) = (1 - s(t))H_I + s(t)H_P \quad (1)$$

The scheduling function is often set as $s(t) = t/T$, where T is a total evolution time length. By using $|\Psi_{\text{HF}}\rangle$ and a Fock operator as the initial wave function and Hamiltonian, respectively, and the full Hamiltonian as H_P , we can obtain $|\Psi_{\text{full-CI}}\rangle$ if the evolution time T is long enough⁴. Note that an approach using the maximum commuting Hamiltonian as the initial Hamiltonian H_I was proposed recently²². ASP was adopted for the QPE-based full-CI of H_2 molecule using an NMR quantum computer²³. ASP numerical simulations of the 1^1A_1 state of methylene (CH_2) molecule was reported in 2014²⁴, and the optimization of the scheduling procedure by using VQE was discussed in 2021²⁵. Application of the complete active space configuration interaction

Table 1 Scheduling functions $s(x)$ ($x = m/M$) tested in Eq. 1.

Name and abbreviation	Function
Linear (Lin)	$s(x) = x$
Square (Squ)	$s(x) = 3x^2 - 2x^3$
Sinusoidal (Sin)	$s(x) = \sin(\pi x/2)$
Sinusoidal cubic (SinCub)	$s(x) = \sin^3(\pi x/2)$
Cubic (Cub)	$s(x) = 6x^5 - 15x^4 + 10x^3$

(CASCI) wave function as the initial wave function and the nonlinear scheduling functions $s(t) = (t/T)^c$ ($0 < c < 1$) was also reported in 2021²⁶.

Although ASP is a long-term (or FCTC) algorithm because the quantum circuit for ASP is usually too deep to execute on NISQ devices, it is promising because it does not require detailed a priori knowledge of the electronic structure of the system being studied. However, ASP is potentially unsuitable for strongly correlated systems, because the evolution time length T should be set longer when it is applied to the systems with smaller energy gaps between the ground and excited states. We also emphasize that the number of ASP studies so far documented are very few, and there has been little knowledge on the optimal computational conditions of ASP, such as the setting of the evolution time length and the selection of scheduling functions. Then, it is essential to explore suitable computational conditions which help make ASP as a practical tool for the preparation of correlated wave functions, which is the main subject of this work. Note that variational quantum imaginary time evolution (QITE) is also available for the preparation of correlated wave functions^{27–30}. However, the total length of imaginary time propagation is determined by the spectrum of the Hamiltonian and the initial overlap with the exact wave function, and the application of QITE to strongly correlated systems is also an important issue.

In this work, we have carried out numerical simulations of ASP for the generation of correlated wave functions in the triple bond dissociation of N_2 molecule, the symmetric bond dissociation of BeH_2 molecule, and the C_{2v} quasi-reaction pathway of beryllium atom insertion to H_2 molecule. We have explored the effects of nonlinear scheduling functions and the ASP starting with a broken-symmetry (BS) wave function $|\Psi_{BS}\rangle$ by using the electron spin S^2 operator as the penalty term in the time-dependent Hamiltonian. As the scheduling function $s(t)$ in Eq. 1, we examined five different functions listed in Table 1 and plotted in Fig. 1; all of them were studied as the scheduling function in adiabatic algorithms by Hu and Wu³¹. An anticipated application of ASP is the wave function preparation for QPE, and thus in this study the quality of the wave functions obtained from ASP is evaluated by means of the square overlap with the CASCI wave function, $|\langle \Psi_{ASP} | \Psi_{CASCI} \rangle|^2$. Note that to use ASP for the wave function preparation in QPE, the evolution time required for ASP must be significantly shorter than that needed in QPE ($T \sim 2000$ atomic unit to achieve 1 kcal mol⁻¹ of energy precision¹⁶). Eventually, we attempt to propose guidelines for the evolution time length and the selection for starting wave functions toward the practical use of ASP.

Results and discussion

Scheduling function and evolution time length dependences of ASP in the potential energy curve of N_2 . First, we explored the scheduling function and the evolution time length dependences of ASP in N_2 molecule with $R(N-N) = 1.0, 2.0,$ and 3.0 Å. These geometries are selected as the representatives of the structures close to the equilibrium geometry where $|\langle \Psi_{HF} | \Psi_{CASCI} \rangle|^2$ is large, in the intermediate bond dissociation region where neither the

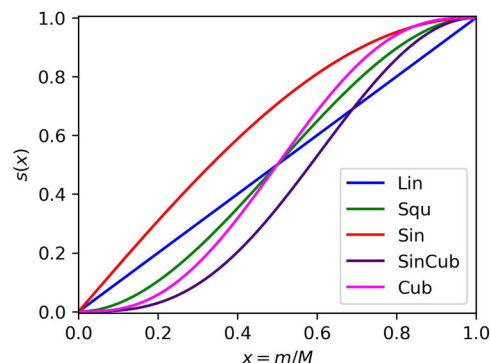


Fig. 1 Plots of the scheduling functions tested in Eq. 1. Mathematical definitions of the scheduling functions are provided in Table 1.

$|\Psi_{HF}\rangle$ nor the $|\Psi_{CSF}\rangle$ are good approximation of the ground state, and in the bond dissociated region where the $|\Psi_{CASCI}\rangle$ can be approximate to a $|\Psi_{CSF}\rangle$. The numerical quantum circuit simulation results with $T = 10$ – 100 are summarized in Fig. 2. The number of quantum gates required for a single Trotter step is about 6900. At the geometry $R(N-N) = 1.0$ Å, ASP gives the wave function close to the $|\Psi_{CASCI}\rangle$ even for the shortest evolution time length being tested ($T = 10$) regardless of the choice of the scheduling functions. The scheduling function and evolution time length dependences become significant for the longer N–N distances. Among the five scheduling functions being tested, the sinusoidal function exhibits the fastest convergence to the CASCI wave function against the evolution time length T . By employing the sinusoidal function as the scheduling function, we need $T = 50$ and 70 for $R(N-N) = 2.0$ and 3.0 Å, respectively, to achieve $|\langle \Psi_{ASP} | \Psi_{CASCI} \rangle|^2 > 0.9$. Longer evolution time is required to achieve the same magnitude of the square overlap for the elongation of the N–N bond, because the HOMO–LUMO gap and the energy gap between the ground and excited states become smaller for the elongated N–N distances. Plots of the S_1 – S_0 energy gap of the instantaneous Hamiltonian (Fig. 3) indicate that the $s(x)$ value giving the minimum $\Delta E(S_1-S_0)$ value becomes larger for the shorter $R(N-N)$ values.

The trajectories of ASP of N_2 in the geometry $R(N-N) = 3.0$ Å with the evolution time length $T = 100$ are plotted in Fig. 4. Among the five scheduling functions, the sinusoidal function gives the smallest $\Delta E_{ASP-CASCI}$ value and the largest square overlap at each time step. Note that the trajectories of the square overlap calculated by using the linear and sinusoidal functions oscillate in the beginning of ASP. This originates from the fact that the gradient $\partial E(t)/\partial s(t) \neq 0$ at $t = 0$ ³¹. The same trends were also observed in the other geometries and molecules under study.

The reason why the sinusoidal function gave the best results among the five scheduling functions can be explained by the structure of the HF wave function and the energy landscape of the instantaneous Hamiltonian. One has to set the evolution time length prior to computation to apply ASP. The evolution time length T is usually set to satisfy the condition in Eq. 2³².

$$\max_{s \in [0,1]} \frac{|\langle \Psi_g(s) | \partial_s H(s) | \Psi_e(s) \rangle|}{|\epsilon_g(s) - \epsilon_e(s)|^2} \ll T \quad (2)$$

Here, $|\Psi_g(s)\rangle$ and $|\Psi_e(s)\rangle$ are the wave functions of the electronic ground and excited states, respectively, of the instantaneous Hamiltonian $H(s)$, and $\epsilon_g(s)$ and $\epsilon_e(s)$ are the corresponding energy eigenvalues. From Eq. 2, we can sweep the time-dependent Hamiltonian faster if the energy gaps between the ground and the excited states are large and if the numerator of Eq. (2) is small.

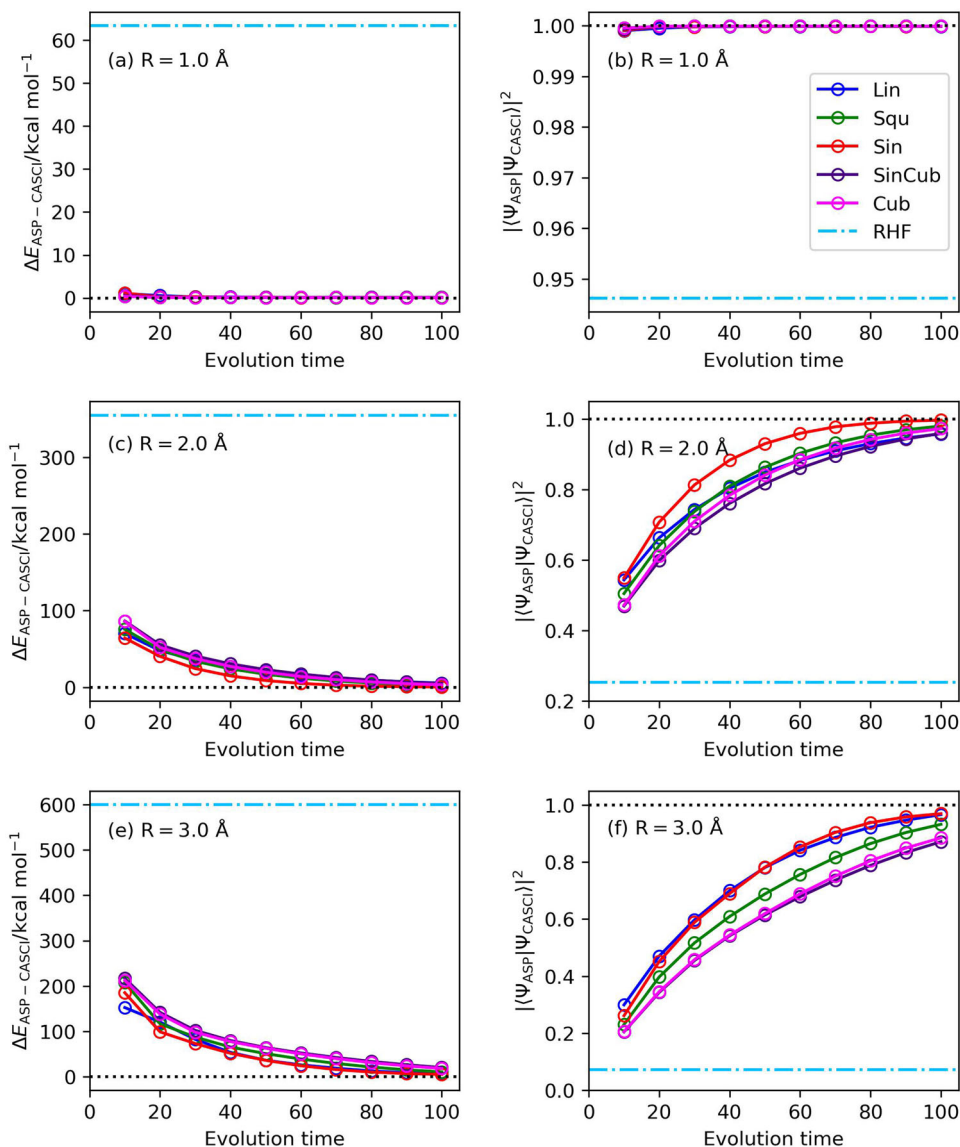


Fig. 2 Results of the numerical quantum circuit simulation of N_2 molecule. **a, c, e** The energy deviations from the CASCI values in **(a)** $R(N-N) = 1.0 \text{ \AA}$, **(c)** $R(N-N) = 2.0 \text{ \AA}$, and **(e)** $R(N-N) = 3.0 \text{ \AA}$. **b, d, f** The square overlaps with the CASCI wave function in **(b)** $R(N-N) = 1.0 \text{ \AA}$, **(d)** $R(N-N) = 2.0 \text{ \AA}$, and **(f)** $R(N-N) = 3.0 \text{ \AA}$.

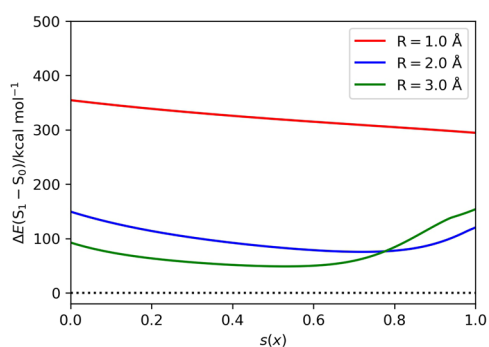


Fig. 3 The S_1-S_0 energy gap of the instantaneous Hamiltonian $H(s(x))$ of N_2 molecule with the bond lengths $R(N-N) = 1.0, 2.0,$ and 3.0 \AA . The $\Delta E(S_1 - S_0)$ values at $s(x) = 0.0$ and 1.0 corresponds to the energy gap of the Fock operator and the full electronic Hamiltonian, respectively.

The $|\Psi_{HF}\rangle$ can be rewritten as $|\Psi_{HF}\rangle = \sum_j c_j |\Psi_j\rangle$, where $|\Psi_j\rangle$ is the j th eigenfunction and c_j is the corresponding coefficient. Because the $|\Psi_{HF}\rangle$ is spin and spatial symmetry-adapted, only the excited states belonging to the same spin and spatial symmetry can have non-negligible contribution to the numerator of Eq. 2 in the absence of Trotter decomposition errors and noises. The excitation energies of the corresponding excited states in the weakly correlated regime are generally larger in the earlier stage of ASP. This is because the Fock operator used as the initial Hamiltonian contains the terms with occupied orbitals only, and the excited states cannot be sufficiently stabilized under the Fock operator. Thus, we can sweep the Hamiltonian faster in the earlier stage of ASP, and the sweep speed must be attenuated by the time evolution. The sinusoidal function has such a structure.

Note that Eq. 2 is insufficient, and it does not guarantee the validity of the adiabatic approximation^{33,34}. In fact, Marzlin and Sanders claimed that the application of the adiabatic theorem may lead to an inconsistency no matter how slowly the Hamiltonian is

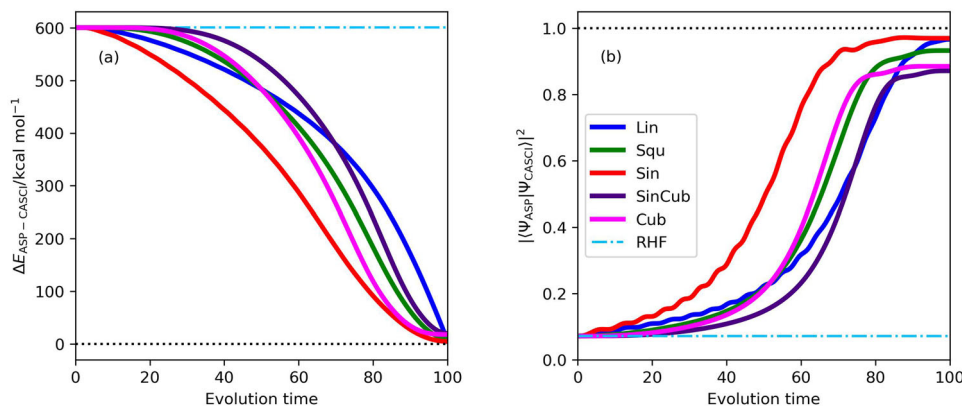


Fig. 4 Trajectories of ASP with $T = 100$ in N_2 molecule with $R(N-N) = 3.0 \text{ \AA}$. **a** The energy deviations from the CASCI value. **b** The square overlaps with the CASCI wave function.

varied³³. They also pointed out that the inconsistency becomes a potential problem whenever $|\Psi(T)\rangle$ deviates greatly from the initial state $|\Psi(0)\rangle$. Strongly correlated systems can be the case of such a small overlap $|\langle \Psi(0) | \Psi(T) \rangle|$ if the $|\Psi_{HF}\rangle$ is employed as $|\Psi(0)\rangle$, and therefore the application of ASP to strongly correlated systems is a challenging problem. The necessary and sufficient condition for adiabatic evolution was discussed by Wang and Plenio, by decomposing the diabatic propagator into the geometric functions determined by the eigenstates and the modulation functions determined by the energy gaps and the speed of sweeping³⁵. By utilizing these necessary and sufficient conditions for adiabatic condition, Xu and coworkers demonstrated the adiabatic evolution in the presence of vanishing energy gaps, using a nitrogen-vacancy center in diamond³⁶. It should be also noted that approaches based on the shortcuts to adiabaticity (STA) have been eagerly studied to manipulate the quantum system on timescales shorter than decoherence time³⁷, but it requires some non-physical Hamiltonian in order to make it work, and it is still an open and challenging problem.

Although Eq. 2 may lead to an inconsistency of the adiabatic evolution^{33,34}, it can be used as the guideline of the evolution time length. Indeed, the variation of the wave function under the time-dependent Hamiltonian, which is responsible for the inconsistency, may be difficult to estimate in advance of the ASP simulation, but the energy gap between the ground and the first excited states can be roughly evaluated from the HOMO–LUMO gap $\Delta\epsilon = \epsilon(\text{LUMO}) - \epsilon(\text{HOMO})$, where $\epsilon(\text{HOMO})$ and $\epsilon(\text{LUMO})$ are the orbital energies of HOMO and LUMO, respectively. We note that the HOMO–LUMO gap estimation is only a practical approach. The exact energy landscape is required to fully adjust the scheduling function $s(x)$, which corresponds to solving the problem altogether. However, the HOMO–LUMO gap can be used for crude estimation of the excitation energy. These orbital energies can be obtained from the HF calculations, and therefore using the HOMO–LUMO gap for the determination of the evolution time length does not need additional computation. Excitation energy estimations based on the single excitation CI or time-dependent DFT may be more reliable, but such computations inherently raise the computational cost. Based on this strategy, we examined the following evolution time determination methods: $T = 5/\Delta\epsilon^2$ and $T = 10/\Delta\epsilon^2$. In addition to them, we also tested the evolution time length $T = 20/\Delta\epsilon$. The results of the quantum circuit simulations with different evolution time length determination strategies with the sinusoidal scheduling function are summarized in Fig. 5. If we adopt the strategy $T = 10/\Delta\epsilon^2$, we can obtain the correlated wave function with the square overlap $|\langle \Psi_{ASP} | \Psi_{CASCI} \rangle|^2 > 0.99$ at all bond lengths under study, but the evolution time for the longer N–N distance is quite long. The

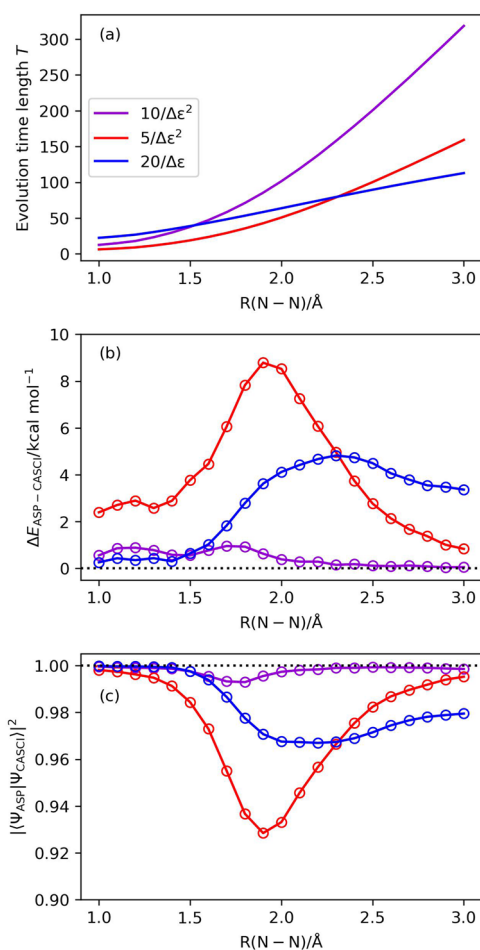


Fig. 5 Results of the numerical simulation of ASP with different evolution time length determination strategies in N_2 . Sinusoidal function is used for the scheduling function. **a** Evolution time length being tested. **b** The energy deviations from the CASCI values. **c** The square overlaps with the CASCI wave functions.

strategy $T = 5/\Delta\epsilon^2$ gives fairly good results except for intermediate bond dissociation region. Considering the evolution time lengths and the square overlap, the strategy $T = 20/\Delta\epsilon$ seems to be most suitable for ASP of the triple bond dissociation in N_2 . The same trend was observed for the other scheduling functions, although the deviations from the CASCI result are larger for the other functions than the sinusoidal one (see Supplementary Fig. S2).

ASP with broken-symmetry wave functions and the S^2 penalty term in the potential energy curve of N_2 . It is naturally expected that the use of multiconfigurational wave functions gives a plausible choice for the starting wave function in strongly correlated systems. In fact, Kremenetski and coworkers reported that substantial speedup of ASP can be achieved by using the CASCI wave function as the starting wave function²⁶. However, encoding the CASCI wave function on a quantum register becomes non-trivial when the active space of CASCI is large. Here, we examine an alternative and facile approach of ASP for strongly correlated systems by adopting the BS wave function $|\Psi_{BS}\rangle$ as the starting wave function with the S^2 operator as the penalty term in the time-dependent Hamiltonian. The BS methods have been widely used in the DFT calculations of open-shell low-spin states such as spin-singlet states of biradicals^{38–40}. The $|\Psi_{BS}\rangle$ is a single Slater determinant with spin- β unpaired electrons in the localized singly occupied molecular orbitals (SOMOs). The $|\Psi_{BS}\rangle$ is an eigenfunction of the S_z operator but is not an eigenfunction of the S^2 operator, and thus the $|\Psi_{BS}\rangle$ is expressed by a linear combination of wave functions having different spin quantum numbers S , as in Eqs. 3 and 4 for four- and six-spin $M_S = 0$ BS wave functions, respectively, for example.

$$|\alpha\alpha\beta\beta\rangle = \frac{1}{\sqrt{6}}|\Psi_{S=2, M_S=0}\rangle + \frac{1}{\sqrt{2}}|\Psi_{S=1, M_S=0}\rangle + \frac{1}{\sqrt{3}}|\Psi_{S=0, M_S=0}\rangle \quad (3)$$

$$|\alpha\alpha\alpha\beta\beta\rangle = \frac{1}{\sqrt{20}}|\Psi_{S=3, M_S=0}\rangle + \frac{1}{2}|\Psi_{S=2, M_S=0}\rangle + \frac{3}{\sqrt{20}}|\Psi_{S=1, M_S=0}\rangle + \frac{1}{2}|\Psi_{S=0, M_S=0}\rangle \quad (4)$$

The coefficients in the right-hand side of Eqs. 3 and 4 can be derived from the structure of spin eigenfunctions⁴¹. The eigenvalue of the S^2 operator is $S(S + 1)$, and the expectation values of the S^2 operator of the wave functions given in Eqs. 3 and 4 are calculated to be 2.0 and 3.0, respectively.

Because the quantum state corresponding to the $|\Psi_{BS}\rangle$ can be prepared on a quantum computer with the same cost as the preparation of $|\Psi_{HF}\rangle$ regardless of the number of singly occupied orbitals, it is possible to use the $|\Psi_{BS}\rangle$ as the starting wave function in ASP. The wave function components having the spin quantum number different from the target electronic state can be eliminated by introducing the S^2 operator as the penalty term in the time-dependent Hamiltonian $H(t)$, as in Eq. 5.

$$H(t) = (1 - s(t))H_I + s(t)H_p + s(t)cS^2 \quad (5)$$

Here, c is a coefficient that controls the strength of the penalty term. The S^2 operator as the penalty term works to raise the energy of the wave function with the spin quantum number S by $cS(S + 1)$, and therefore spin contaminants can be readily eliminated during ASP. We expect that ASP starting from $|\Psi_{BS}\rangle$ can generate multiconfigurational wave functions efficiently, just as the spin-projected extended Hartree-Fock (EHF) method⁴² in classical computation that applies the spin projection operator to the spin contaminated UHF wave function. In Eq. 3, larger c values can shift the energies of the spin contaminants greatly, but too large c values will result in departure from the adiabatic pathway. Note that we have already proposed a method to construct a quantum circuit for the time evolution operator $\exp(-is(t)cS^2t)$ by utilizing a generalized spin coordinate mapping⁴³, which can be directly used for the present ASP study.

The results of the ASP starting from the six-spin $|\Psi_{BS}\rangle$ (the wave function in Eq. 4) with $c = 0.5$, the square scheduling function, and the evolution time length $T = 10$ –100 of N_2

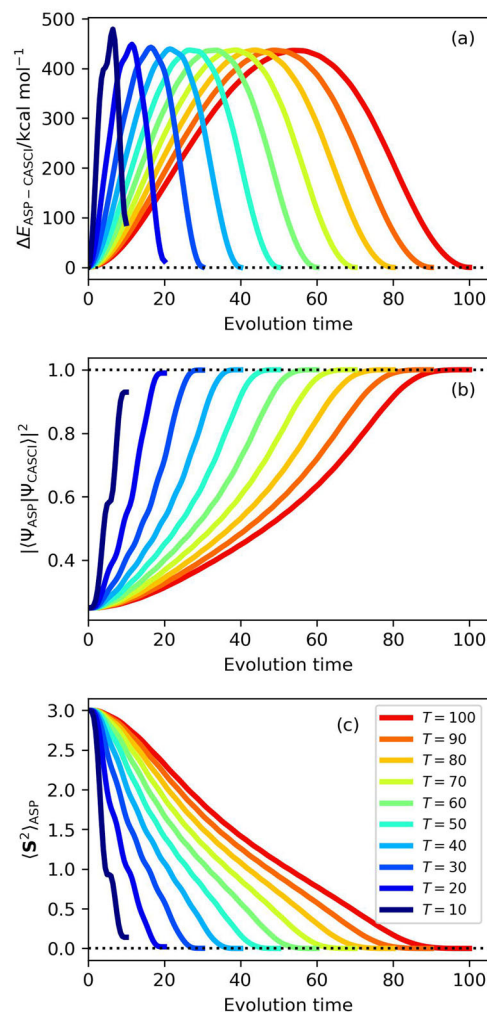


Fig. 6 Results of the numerical simulation of ASP with $|\Psi_{BS}\rangle$ as the starting wave function and square function as the scheduling function in N_2 molecule at $R(N-N) = 3.0 \text{ \AA}$. **a** The energy deviations from the CASCI values. **b** The square overlaps with the CASCI wave functions. **c** The $\langle S^2 \rangle$ values.

molecule at $R(N-N) = 3.0 \text{ \AA}$ are summarized in Fig. 6, and the results obtained by using the other scheduling functions are given in Supplementary Fig. S3–S6. Interestingly, ASP starting with the $|\Psi_{BS}\rangle$ gave the square overlap $|\langle \Psi_{ASP} | \Psi_{CASCI} \rangle|^2$ larger than 0.995 even for $T = 20$. The scheduling function dependence on ASP with the $|\Psi_{BS}\rangle$ with the total evolution time $T = 50$ were plotted in Fig. 7. The $|\langle \Psi_{ASP} | \Psi_{CASCI} \rangle|^2$ values calculated by using the square, sinusoidal cubic, and cubic scheduling functions asymptotically approach unity for longer N–N distances. These three scheduling functions exhibit the best square overlap values in the bond dissociation region. No significant differences were observed among these scheduling functions, but the square function gave larger square overlap than the sinusoidal cubic and cubic functions for shorter N–N bond lengths. Importantly, the square, sinusoidal cubic, and cubic scheduling functions gave almost spin-pure wave functions with $\langle S^2 \rangle_{ASP} \approx 0$, although spin contaminations were not eliminated completely in the $|\Psi_{ASP}\rangle$ generated by employing the sinusoidal and linear scheduling functions. In the following discussions we used the square scheduling function for ASP with the $|\Psi_{BS}\rangle$ wave functions.

Again, the reason why the square, sinusoidal cubic, and cubic scheduling functions gave good results can be explained by the structure of wave functions and the energy landscape of the

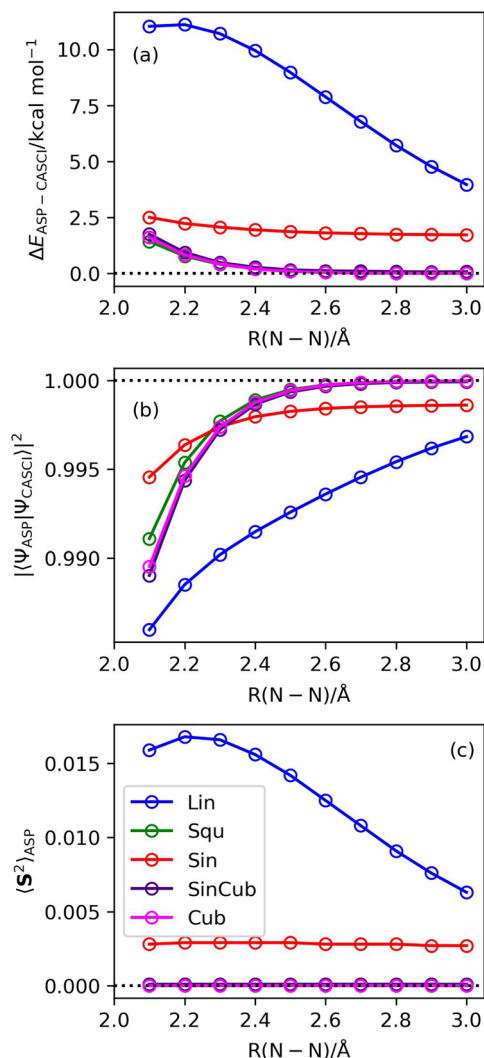


Fig. 7 Numerical simulation results of the scheduling function dependence on ASP with the $|\Psi_{\text{BS}}\rangle$ as the starting wave function and $T = 50$ in N_2 molecule. **a** The energy deviations from the CASCI values. **b** The square overlaps with the CASCI wave functions. **c** The $\langle S^2 \rangle$ values.

adiabatic evolution pathway. As given in Eqs. 3 and 4, the $|\Psi_{\text{BS}}\rangle$ is expressed by the linear combination of electronic states belonging to different spin multiplicities. These electronic states have similar energies in the beginning of ASP, but the quasi-degeneracy is gradually lifted as the ASP proceeds, owing to the S^2 operator introduced as the penalty term in the Hamiltonian. Therefore, variation of the Hamiltonian must slow in the earlier stage of ASP, if the $|\Psi_{\text{BS}}\rangle$ is used as the starting Hamiltonian. All the square, sinusoidal cubic, and cubic functions have zero gradient $ds(x)/dx = 0$ at $x = 0$, which is very important to acquire good wave functions in the ASP with the $|\Psi_{\text{BS}}\rangle$.

Note that ASP with the $|\Psi_{\text{BS}}\rangle$ has not only advantages described above but also disadvantages. The BS method generally breaks the spatial symmetry as well as the spin symmetry, because the localized orbitals are prepared by taking a linear combination of HOMO $-i$ and LUMO $+i$ orbitals. In the ASP starting from $|\Psi_{\text{HF}}\rangle$, we can utilize the point group symmetry to reduce the number of the nonzero Hamiltonian terms, but such reduction is not applicable for the $|\Psi_{\text{BS}}\rangle$ with localized MOs. In addition, the computational cost of the time evolution operator under the S^2 operator, $\exp(-iS^2t)$ also pushes the computational cost of ASP. The scaling of the computational cost of simulating $\exp(-iS^2t)$ is

$O(n_{\text{MO}}^2M)$, where n_{MO} is the number of molecular orbitals in the active space and M refers to the number of ASP steps. As a result, the computational cost of each ASP step is larger by using the $|\Psi_{\text{BS}}\rangle$ than that for using the $|\Psi_{\text{HF}}\rangle$, especially for the molecular systems with high point group symmetry. However, for large molecular systems, it is possible to take advantage of the locality of MOs to reduce the computational cost, by ignoring the Hamiltonian terms having the norms smaller than the threshold⁴⁴. It should be also noted that our numerical simulations suggest QITE can be also accelerated by using the $|\Psi_{\text{BS}}\rangle$ as the starting wave function in conjunction with the S^2 penalty term in the Hamiltonian.

A criterion for the switching of the starting wave function. In the previous section, we demonstrated that ASP using the $|\Psi_{\text{BS}}\rangle$ converges quickly in the region of the bond dissociation. However, we can expect that the $|\Psi_{\text{HF}}\rangle$ is more suitable for the starting wave function of ASP around the equilibrium geometry. It should be also noted that the BS-UHF calculation converges to the RHF solution for shorter bond lengths. Constructing a guideline for the switching of the starting wave function is important toward practical application of ASP to other molecular systems.

Here, we have examined a criterion for selecting the starting wave function based on diradical characters y_i ^{45,46}. Diradical characters are used as the measure of open shell electronic configurations. At the spin-projected UHF level, they can be calculated from the occupation number of the unoccupied natural orbitals $n_{\text{LUNO}+i}$ using Eq. 6⁴⁶.

$$y_i = 1 - \frac{2(1 - n_{\text{LUNO}+i})}{1 + (1 - n_{\text{LUNO}+i})^2} \quad (6)$$

In the triple bond dissociation of N_2 molecule, two types of diradical characters can have significant values: $y(\pi) = y_0 = y_1$ and $y(\sigma) = y_2$, those reflect open shell electronic configurations in the valence π and σ bonds, respectively. Thus, there are two possible choices of the $|\Psi_{\text{BS}}\rangle$ in the ASP study of the potential energy curve of N_2 molecule, $|\Psi_{\text{BS}2}\rangle$ and $|\Psi_{\text{BS}3}\rangle$. The $|\Psi_{\text{BS}2}\rangle$ is the four-spin BS wave function where two π orbitals are treated by means of the BS approach. The $|\Psi_{\text{BS}3}\rangle$ is a six-spin BS wave function and both the σ and π orbitals are dealt with the BS framework.

We assume that the BS treatment is more feasible if the corresponding diradical character is larger. Following this assumption, we explored the threshold value of the diradical characters for the switching of the starting wave function. By setting the evolution time length $T = 20/\Delta\epsilon$ for the ASP with the $|\Psi_{\text{HF}}\rangle$ and $T = 50$ for that with the $|\Psi_{\text{BS}}\rangle$, we found that setting the threshold value for the diradical characters to be 0.6 gives fairly good results, from the viewpoints of both the evolution time length and quality of the $|\Psi_{\text{ASP}}\rangle$, in the potential energy curve of N_2 molecule (Fig. 8). This is the result of particular ASP conditions, N_2 molecule with (6e,6o) active space and STO-3G basis set with the sinusoidal and square scheduling functions for the $|\Psi_{\text{HF}}\rangle$ and the $|\Psi_{\text{BS}}\rangle$, respectively, and it is natural that different molecular systems and different computational conditions will give different optimal threshold values for the switching of the starting wave function. We also carried out the ASP simulations of N_2 molecule using the 6-31G* and 6-311G* basis sets with (6e,6o), (10e,8o), (6e,8o), and (10e,10o) active spaces, obtaining qualitatively the same results (see Supplementary Note 4). Nevertheless, the exploration of other molecular systems is necessary to get further insight of ASP. In the following sections, we describe the results of ASP in the potential energy curve under the symmetric Be-H bond dissociation in the linear

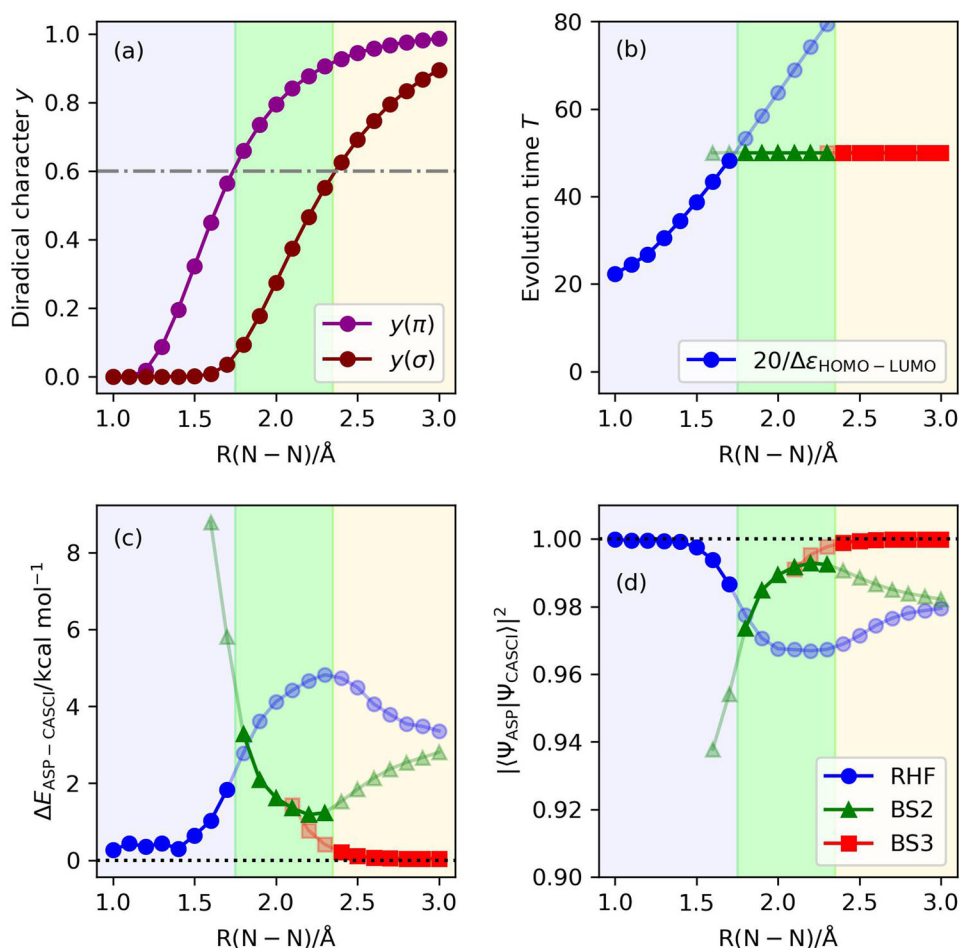


Fig. 8 Results of the numerical simulation of ASP of the potential energy curve for N_2 molecule, using the diradical characters as the indicator for selecting the initial wave function. Background colors specify the regions of the initial wave function recommended from the criterion based on the diradical character $y > 0.6$. Sinusoidal and square functions were adopted as the scheduling functions with the $|\Psi_{HF}\rangle$ and $|\Psi_{BS}\rangle$, respectively, as the starting wave function. **a** Diradical characters calculated using Eq. 6. **b** Evolution time lengths. **c** The energy differences from the CASCI values. **d** The square overlaps with the CASCI wave functions.

BeH_2 molecule, and the C_{2v} quasi-reaction pathway of the $Be + H_2 \rightarrow BeH_2$ reaction.

Potential energy curve of the symmetric bond dissociation in the linear BeH_2 . BeH_2 is a linear molecule with $R(Be-H) = 1.326$ Å in the equilibrium geometry⁴⁷. Symmetric elongation of two $Be-H$ bonds generates a Be atom in the $(1s)^2(2s)^2$ closed-shell singlet electron configuration and two H atoms. In this work, we have explored the potential energy curve in the range of $R(Be-H)$ from 0.7 to 4.0 Å. The BS-UHF/STO-3G calculations converged to the RHF solution at the bond length $R(Be-H) = 1.9$ Å and shorter. The calculated diradical character y is plotted in Fig. 9a. By adopting the criterion for selecting the starting wave function discussed in the previous section, we expect that the $|\Psi_{BS}\rangle$ is suitable for the starting wave function of ASP at the bond length $R(Be-H) = 2.5$ Å and longer. The quantum circuit simulation results are summarized in Fig. 9. By switching the starting wave function from the $|\Psi_{HF}\rangle$ to the $|\Psi_{BS}\rangle$ at the bond length $R(Be-H) = 2.5$ Å, we obtained the $|\Psi_{ASP}\rangle$ with the square overlap $|\langle\Psi_{ASP}|\Psi_{CASCI}\rangle|^2 > 0.98$ for all the bond lengths being studied. These results exemplify the usefulness of the diradical character as the indicator of the starting wave function switching. However, the $|\langle\Psi_{ASP}|\Psi_{CASCI}\rangle|^2$ values are slightly larger in the geometries with intermediate diradical characters ($R(Be-H) \sim 2.4$ Å). The ASP simulations with the longer evolution time revealed that we can achieve $|\langle\Psi_{ASP}|\Psi_{CASCI}\rangle|^2 > 0.998$ for $T = 200$ and 100

with $|\Psi_{HF}\rangle$ and $|\Psi_{BS}\rangle$, respectively, as the starting wave function (see Supplementary Figs. S16 and S17). We also examined the numerical simulations by using the other scheduling functions for ASP with the $|\Psi_{HF}\rangle$ and the evolution time length $T = 20/\Delta\epsilon$. The results are summarized in Supplementary Fig. S18, insisting that the sinusoidal function is suitable for the scheduling function.

The C_{2v} quasi-reaction pathway of Be atom insertion to H_2 . Finally, we have examined ASP in the C_{2v} quasi-reaction pathway of the Be atom insertion to H_2 molecule. This system has been widely studied as the model of strongly correlated electronic structures^{48–52}. Cartesian coordinates of H atoms are summarized in Table 2. The reaction pathway contains the S_0-S_1 avoided crossing at the transition structure (point E in Table 2), and the energy gap between the S_0 and S_1 states becomes small around the transition structure.

Results of the numerical quantum circuit simulations are summarized in Fig. 10. By using the $|\Psi_{HF}\rangle$ as the starting wave function and setting the evolution time length as $T = 20/\Delta\epsilon$ and using the sinusoidal scheduling function, we obtained the correlated wave function with the square overlap close to unity, except for point E. To disclose the convergence behavior of ASP at point E, we examined the ASP simulations with longer evolution time lengths $T = 200, 300, 400,$ and 500 . The square overlaps were calculated to be $|\langle\Psi_{ASP}|\Psi_{CASCI}\rangle|^2 = 0.9308, 0.9720,$

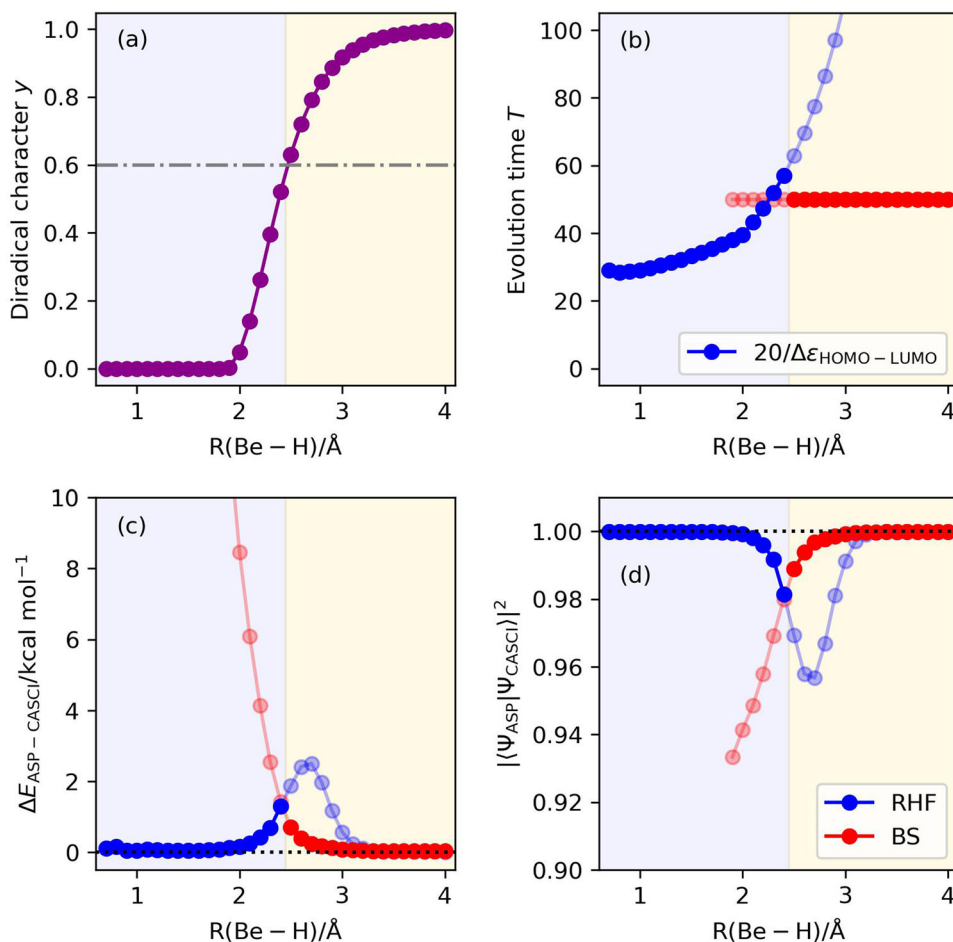


Fig. 9 Results of the numerical simulation of ASP of the potential energy curve of the symmetric bond dissociation in BeH₂ molecule, using the diradical character as the indicator for selecting the initial wave function selection. Background colors specify the regions of the initial wave function recommended from the criterion based on the diradical character $y > 0.6$. Sinusoidal and square functions were adopted as the scheduling functions with the $|\Psi_{\text{HF}}\rangle$ and $|\Psi_{\text{BS}}\rangle$, respectively, as the starting wave function. **a** Diradical characters calculated using Eq. 6. **b** Evolution time lengths. **c** The energy differences from the CASCI values. **d** The square overlaps with the CASCI wave functions.

Table 2 Cartesian coordinates of H atoms for the points being studied in the Be + H₂ → BeH₂ reaction, in units of Bohr^[a].

Point	X	Y	Z
A	0.000	±2.540	0.000
B	0.000	±2.080	1.000
C	0.000	±1.620	2.000
D	0.000	±1.390	2.500
E	0.000	±1.275	2.750
F	0.000	±1.160	3.000
G	0.000	±0.930	3.500
H	0.000	±0.700	4.000
I	0.000	±0.700	6.000

^[a]Be atom is located at the origin of coordinates.

0.9894, and 0.9962, respectively, and thus the convergence of the square overlap against the evolution time length T is very slow. The simulations around point E with finer geometrical changes revealed that the square overlap is smaller for the geometry closer to the transition structure (see Supplementary Note 7). We also computed the energy landscape of the instantaneous Hamiltonian at point E, finding that the S_1 and S_0 states become almost gapless

around $s(x) = 0.9$ (see Supplementary Fig. S19). The 16, 14, and 12 qubit ASP simulations by removing the highest virtual orbitals one by one gave the square overlaps $|\langle \Psi_{\text{ASP}} | \Psi_{\text{CASCI}} \rangle|^2 = 0.8464$, 0.8538, and 0.9751, respectively. These results exemplify that selecting appropriate active orbitals is essential to obtain sophisticated wave functions from ASP. In fact, 16 qubit ASP simulations using the natural orbitals constructed from CISD/6-31G* calculations gave the square overlap larger than 0.98 for all points being studied (see Supplementary Fig. S21).

The BS-UHF calculations revealed that points D, E, and F have non-negligible diradical characters. The diradical characters were calculated to be $y = 0.2991$, 0.7851, and 0.6125 for points D, E, and F, respectively. The diradical character indicates that the $|\Psi_{\text{BS}}\rangle$ is suitable for the starting wave function of ASP in points E and F. Our numerical simulations revealed that ASP with the $|\Psi_{\text{BS}}\rangle$ gave larger square overlap values with the CASCI wave functions not only at points E and F but also at point D, with shorter evolution time length than ASP with the $|\Psi_{\text{HF}}\rangle$. More extended studies are necessary to thoroughly understand the performance of ASP, which is left as future work.

Conclusions

In this work, we have examined the numerical quantum circuit simulations of ASP in N₂ and BeH₂ molecules, seeking practical computational conditions for the generation of correlated wave

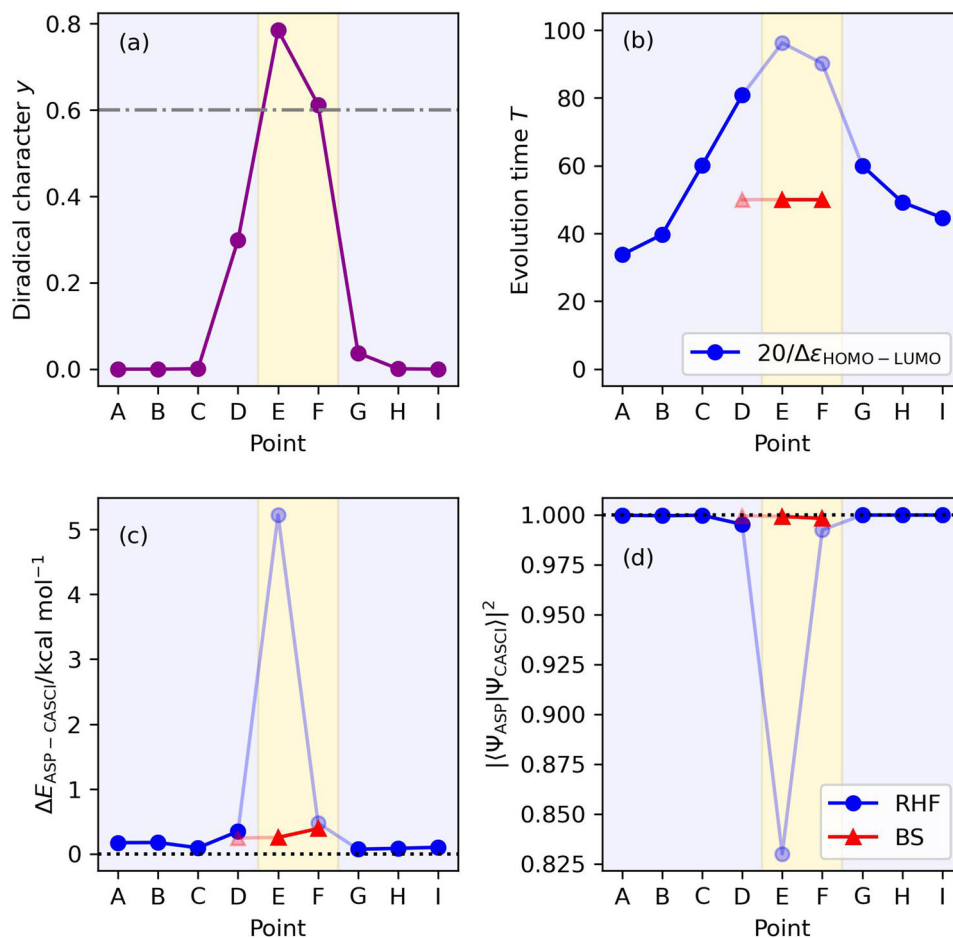


Fig. 10 Results of the numerical simulation of ASP of the C_{2v} quasi-reaction pathway of $\text{Be} + \text{H}_2 \rightarrow \text{BeH}_2$, using the diradical character as the indicator for selecting the initial wave function. Background colors specify the regions of the initial wave function recommended from the criterion based on the diradical character $\gamma > 0.6$. Sinusoidal and square functions were adopted as the scheduling functions with the $|\Psi_{\text{HF}}\rangle$ and $|\Psi_{\text{BS}}\rangle$, respectively, as the starting wave function. **a** Diradical characters calculated using Eq. 6. **b** Evolution time lengths. **c** The energy differences from the CASCI values. **d** The square overlaps with the CASCI wave functions.

functions having the square overlap with the CASCI wave function close to unity. The numerical quantum circuit simulations revealed that if the $|\Psi_{\text{HF}}\rangle$ is employed as the starting wave function, the sinusoidal function $s(x) = \sin(\pi x/2)$, where $x = t/T$, gives the wave function having large square overlaps for shorter evolution time lengths among the five scheduling functions being tested. By using the $|\Psi_{\text{HF}}\rangle$ as the starting wave function, the evolution time length required to achieve sufficiently large square overlap with the CASCI wave function increases with increasing the open-shell character and decreasing the HOMO–LUMO gap. By setting the evolution time length $T = 20/\Delta\epsilon$, where $\Delta\epsilon$ denotes the HOMO–LUMO gap, we can obtain the correlated wave function with the square overlap larger than 0.95, except for point E as given in the $\text{Be} + \text{H}_2 \rightarrow \text{BeH}_2$ reaction pathway. By contrast, by using the $|\Psi_{\text{BS}}\rangle$ as the starting wave function and introducing the S^2 operator as the penalty term in the time-dependent Hamiltonian, the ASP simulations with the square function $s(x) = 3x^2 - 2x^3$, sinusoidal cubic function $s(x) = \sin^3(\pi x/2)$, and cubic function $s(x) = 6x^5 - 15x^4 + 10x^3$ gave the correlated wave function with the square overlap close to unity with substantially shorter evolution time length than ASP with the $|\Psi_{\text{HF}}\rangle$ when the diradical character γ is large. The results of the present numerical quantum circuit simulations indicate that ASP is capable of generating the correlated wave functions with sufficiently large square overlap with the CASCI wave function by switching the

starting wave function from the $|\Psi_{\text{HF}}\rangle$ to the $|\Psi_{\text{BS}}\rangle$ and simultaneously changing the scheduling function from the sinusoidal to the square functions, when the diradical character exceeds 0.6. ASP numerical simulations with larger basis sets revealed the importance of the appropriate active space selections based on the occupation number of natural orbitals to acquire sophisticated wave functions from ASP.

It should be emphasized that the computational conditions described in this paper do not have to be optimal for any molecular systems. The number of theoretical and experimental studies of ASP is not many and more elaborated investigations are necessary to shed light on the performance of ASP. Another important direction of the study of ASP is to connect to the QPE algorithms including the Bayesian phase difference estimation algorithm, which is a controlled-time evolution-free algorithm^{5,53}. The relevant studies are underway and will be published in the forthcoming paper.

Methods

Adiabatic state preparation. Assume that the quantum system is in the ground state of an initial Hamiltonian H_I at $t = 0$, and the Hamiltonian of the system changes slowly. If the variation of the Hamiltonian is sufficiently slow, the system remains in the ground state of the instantaneous Hamiltonian at $t > 0$, which is known as an adiabatic theorem¹⁹. ASP utilizes the adiabatic theorem to obtain the full-CI wave function, by using the Fock operator F defined in Eq. 7 as H_I and the

electronic Hamiltonian H in Eq. 8 as H_p .

$$F = \sum_i h_{ii} a_i^\dagger a_i + \frac{1}{2} \sum_{ij} \left(h_{ijij} a_i^\dagger a_j^\dagger a_j a_i + h_{ijij} a_i^\dagger a_j^\dagger a_i a_j \right) \quad (7)$$

$$H = \sum_{pq} h_{pq} a_p^\dagger a_q + \frac{1}{2} \sum_{pqrs} h_{pqrs} a_p^\dagger a_q^\dagger a_s a_r \quad (8)$$

Throughout this paper, we used indices i and j for the occupied spin orbitals in the starting wave function, a and b for the unoccupied spin orbitals, and p, q, r , and s for general spin orbitals. h_{pq} and h_{pqrs} are one- and two-electron molecular orbital (MO) integrals defined in Eqs. 9 and 10, respectively. a_p^\dagger and a_p are creation and annihilation operators, respectively, acting on the p th spin orbital.

$$h_{pq} = \int \phi_p^*(\mathbf{r}) \left(-\frac{1}{2} \nabla^2 - \sum_A \frac{Z_A}{|\mathbf{r} - \mathbf{R}_A|} \right) \phi_q(\mathbf{r}) d\mathbf{r} \quad (9)$$

$$h_{pqrs} = \iint \phi_p^*(\mathbf{r}) \phi_q^*(\mathbf{r}') \frac{1}{|\mathbf{r} - \mathbf{r}'|} \phi_r(\mathbf{r}') \phi_s(\mathbf{r}) d\mathbf{r} d\mathbf{r}' \quad (10)$$

In Eq. 9, A runs over atoms, and Z_A and \mathbf{R}_A are the atomic number and spatial coordinates, respectively, of atom A . ϕ_p is the spatial part of the spin orbital of p .

In order to implement ASP on a gate model quantum computer, the adiabatic evolution is usually discretized: the evolution time length T is divided into M steps, and the time evolution under the time-independent Hamiltonian H_m in Eqs. 11 and 12 is simulated.

$$|\Psi(T)\rangle = e^{-iH_M T/M} \dots e^{-iH_2 T/M} \cdot e^{-iH_1 T/M} |\Psi_1\rangle \quad (11)$$

$$H_m = (1 - s(m/M)) H_I + s(m/M) H_p \quad (12)$$

There are several error sources in ASP. For example, if the evolution time length T is too short to follow the adiabatic path or if the step number M is too small and the Hamiltonian of the $(m+1)$ th step is too different from that of the m th step, the probability to cause nonadiabatic transitions to other electronic states becomes large and the wave function obtained from ASP is expected to have smaller overlap with the exact wave function. Error sources inherent in quantum computing such as decoherence and errors arising from Trotter decomposition, as discussed in the next section, also affect the quality of the wave function. Note that the computational cost of ASP increases linearly to the step number M , although the finer time steps generally give the wave function with the larger overlap with the exact wave function.

Quantum chemical calculations on a quantum computer. To execute ASP on a gate model quantum computer, wave functions are mapped onto qubits by using a fermion–qubit transformation technique, and the quantum circuit corresponding to the time evolution operator is constructed by using basic quantum gate sets⁵⁴. Several fermion–qubit transformation techniques have been proposed^{55–58}, and in this work we adopted a Jordan–Wigner transformation (JWT)^{4,55}. In the JWT, the wave function of the N spin-orbital systems is mapped onto N qubits, and each qubit stores an occupation number of a particular spin orbital: the qubit is in the $|1\rangle$ state if the corresponding spin orbital is occupied by an electron, otherwise in the $|0\rangle$ state. The creation and annihilation operators appearing in the second quantized Hamiltonian (Eqs. 7 and 8) are transformed onto the direct products of Pauli operators (Pauli strings) using Eqs. 13 and 14, respectively.

$$a_p^\dagger = \frac{1}{2} (X_p - iY_p) \otimes \prod_{r=1}^{p-1} Z_r \quad (13)$$

$$a_p = \frac{1}{2} (X_p + iY_p) \otimes \prod_{r=1}^{p-1} Z_r \quad (14)$$

Here, X_p , Y_p , and Z_p are Pauli operators defined in Eqs. 15–17, acting on the p th qubit.

$$X = \begin{pmatrix} 0 & 1 \\ 1 & 0 \end{pmatrix} \quad (15)$$

$$Y = \begin{pmatrix} 0 & -i \\ i & 0 \end{pmatrix} \quad (16)$$

$$Z = \begin{pmatrix} 1 & 0 \\ 0 & -1 \end{pmatrix} \quad (17)$$

By applying the JWT, the second quantized Hamiltonians in Eqs. 7 and 8 are transformed onto qubit Hamiltonians consisting of a linear combination of Pauli strings, as in Eqs. 18 and 19.

$$H = \sum_{k=1}^K \omega_k P_k \quad (18)$$

$$P_k = \sigma_N \otimes \sigma_{N-1} \otimes \dots \otimes \sigma_1, \sigma \in \{I, X, Y, Z\} \quad (19)$$

The time evolution operator U is defined as in Eq. 20. Trotter–Suzuki decomposition^{59,60} is usually used to decompose the time evolution operator and to construct the corresponding quantum circuits. The time evolution operators

obtained by applying the first- and second-order Trotter–Suzuki decompositions are given in Eqs. 21 and 22, respectively⁶¹.

$$U = \exp(-iHt) = \exp(-i \sum_{k=1}^K \omega_k P_k t) \quad (20)$$

$$U \approx \left[\prod_{k=1}^K \exp(-i\omega_k P_k t/L) \right]^L \quad (21)$$

$$U \approx \left[\prod_{k=1}^K \exp(-i\omega_k P_k t/2L) \prod_{k=K}^1 \exp(-i\omega_k P_k t/2L) \right]^L \quad (22)$$

Note that magnitude of the Trotter decomposition error depends on the ordering of terms. It is known that the magnitude ordering in which Hamiltonian terms are applied in the descending order of the absolute value of the coefficient $|\omega_k|$ often gives smaller Trotter decomposition errors than the lexicographical ordering that is an ordering scheme, which maximizes the similarity of the Pauli strings of adjacent terms^{62,63}. Once the Trotter decomposition is applied, the quantum circuit corresponding to the operator $\exp(-i\omega_k P_k t/L)$ is constructed by following the literature³⁴. Supplementary Fig. S1 illustrates the quantum circuit corresponding to the operator $\exp(-i\omega X_0 Z_1 Z_2 X_3 t)$ as an example. Definitions of the quantum gates are also given in the Supplementary Note 1.

Implementation of ASP. ASP can be implemented by the following procedures. (1) Perform the RHF or BS-UHF calculations using conventional quantum chemistry program packages and compute the molecular integrals defined in Eqs. 9 and 10. (2) Divide the electronic Hamiltonian (Eq. 8) into the Fock operator (Eq. 7) and the rest terms, and apply fermion–qubit transformation to them to obtain corresponding qubit Hamiltonians. (3) Set the computational conditions for ASP. Select the initial wave function, scheduling function for adiabatic evolution, total evolution time, time for the single Trotter step, strategy for Trotter term ordering, and so on. (4) Compute the instantaneous Hamiltonian at each discretized time step, and construct the quantum circuit corresponding to the time evolution operator. (5) Encode the starting wave function on the N_{orb} of qubits, where N_{orb} is the number of spin orbitals in the active space, and execute the quantum circuit constructed in the step 4.

The steps 1–4 above are performed on a classical computer, and the step 5 is the main part of the ASP computation. If we adopted $|\Psi_{\text{HF}}\rangle$ as the starting wave function and the JWT for fermion–qubit transformation, the wave function encoding described in the step 5 can be done by applying the Pauli-X (NOT) gates to the qubits storing the occupation number of occupied orbitals in the $|\Psi_{\text{HF}}\rangle$ to the quantum states initialized to the $|00\dots 0\rangle$ state.

Note that in most of adiabatic quantum computing problems the initial Hamiltonian contains the terms those are not included in the final Hamiltonian. On the contrary, all the terms in the Fock operator in Eq. 7 are included in the electronic Hamiltonian in Eq. 8. Therefore, the instantaneous Hamiltonian $H(m)$ defined in Eq. 12 can be rewritten as in Eq. 23, where Fock specifies the qubit Hamiltonian corresponding to the Fock operator.

$$H(m) = \sum_{P_k \in \text{Fock}} \omega_k P_k + \sum_{P_l \notin \text{Fock}} s(m/M) \omega_l P_l \quad (23)$$

In the quantum circuit level, the quantum circuit at each time step has exactly the same structure with different rotational angles θ of the R_z gate (see Supplementary Fig. S1).

Computational conditions. In this work, we have focused on three molecular systems: the potential energy curve of the triple bond dissociation of N_2 molecule, the symmetric Be–H bond dissociations in linear BeH_2 molecule, and the C_{2v} quasi-reaction pathway of a Be atom insertion to a H_2 molecule, at the CASCI level of theory. For the study of the potential energy curve of N_2 molecule under the triple bond dissociation, we used the STO-3G basis set in conjunction with the six electrons in the six orbital (6e, 6o) active space consisting of valence σ/σ^* and π/π^* orbitals. The potential energy curve of BeH_2 molecule under the symmetric Be–H bond cleavage was studied by using the STO-3G basis set and the full-valence (4e, 6o) active space. The quasi-reaction pathway of $\text{Be} + \text{H}_2 \rightarrow \text{BeH}_2$ was investigated by using the basis set comprised of (10s 3p)/[3s 1p] for Be and (4s)/[2s] for H, which was used by Purvis and coworkers⁴⁴ for the study of the same system. Frozen core orbital approximation was adopted for the CASCI calculations and thus the active space is (4e, 9o).

For the numerical quantum circuit simulations of ASP executable on classical computers, we developed a python program by utilizing Cirq⁶⁴ and OpenFermion⁶⁵ libraries. The step number M in Eqs. 11 and 12 was set to be $M = \lceil 2T \rceil$ using a ceil function, and the quantum circuit for the time evolution operator of each step was constructed by adopting the second-order Trotter–Suzuki decomposition with $L = 1$ in Eq. 22. We used the magnitude ordering for the Trotterized time evolution operators, and the ordering of the terms were optimized for every time step.

For the preparation of the starting wave functions of ASP, we performed the RHF and the BS-UHF calculations using GAMESS-US program package⁶⁶. One- and two-electron atomic orbital integrals were also computed using GAMESS-US software, and MO integrals h_{pq} and h_{pqrs} in Eqs. 9 and 10 were generated by using our own AO \rightarrow MO integral transformation program.

Data availability

The data that support the findings on this study are available from the corresponding authors on reasonable request.

Code availability

The python source code for the numerical quantum circuit simulations of ASP and integral files are available via github repository: <https://github.com/Kenji-Sugisaki/ASP>.

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Author contributions

K.Su., K.Sa., and T.T. planned and conducted the project. K.Su. developed the numerical simulation programs and performed quantum chemical calculations and numerical simulations. K.T. and D.S. discussed the results. K.Su. and T.T. wrote the paper.

Competing interests

The authors declare no competing interests.

Additional information

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