

Nonconvergence of the Feynman-Dyson diagrammatic perturbation expansion of propagatorsSo Hirata ^{*}*Department of Chemistry, University of Illinois at Urbana-Champaign, Urbana, Illinois 61801, USA*Ireneusz Grabowski *Institute of Physics, Faculty of Physics, Astronomy, and Informatics, Nicolaus Copernicus University in Toruń, ul. Grudziądzka 5, 87-100 Toruń, Poland*J. V. Ortiz *Department of Chemistry and Biochemistry, Auburn University, Auburn, Alabama 36849-5312, USA*

Rodney J. Bartlett

Quantum Theory Project, Departments of Chemistry and Physics, University of Florida, Gainesville, Florida 32611, USA

(Received 1 February 2024; accepted 29 April 2024; published 16 May 2024)

Using a general-order *ab initio* many-body Green's function method, we numerically illustrate several pathological behaviors of the Feynman-Dyson diagrammatic perturbation expansion of one-particle many-body Green's functions as electron Feynman propagators. (i) The perturbation expansion of the frequency-dependent self-energy is not convergent at the exact self-energy in many frequency domains. (ii) An odd-perturbation-order self-energy has a qualitatively wrong shape and, as a result, many roots of the corresponding Dyson equation are nonphysical in that the poles may be complex or residues can exceed unity or be negative. (iii) A higher even-order self-energy consists of vertical lines at many frequencies, predicting numerous phantom poles with zero residues. (iv) Infinite partial resummations of diagrams by vertex or edge renormalization tend to exacerbate these pathologies. (v) The nonconvergence is caused by the nonanalyticity of the rational-function form of the exact Green's function at many frequencies, where the radius of convergence of its Taylor expansion is zero. This is consistent with the fact that (vi) Padé approximants (power-series expansions of a rational function) can largely restore the correct shape and poles of the Green's function. Nevertheless, not only does the nonconvergence render higher-order Feynman-Dyson diagrammatic perturbation theory useless for many lower-lying ionization or higher-lying electron-attachment states but it also calls into question the validity of its combined use with the *Ansätze* requiring the knowledge of all poles and residues. Such *Ansätze* include the Galitskii-Migdal identity, the self-consistent Green's function methods, and some models of the algebraic diagrammatic construction.

DOI: [10.1103/PhysRevA.109.052220](https://doi.org/10.1103/PhysRevA.109.052220)**I. INTRODUCTION**

In an influential paper [1], Dyson argued that the Feynman-Dyson diagrammatic perturbation theory for quantum electrodynamics is inherently divergent in the presence of electron-positron pair formations even after mass and charge are renormalized. In another important paper [2], Kohn and Luttinger predicted that the finite-temperature diagrammatic perturbation theory for electrons [3–7] does not necessarily reduce to the zero-temperature counterpart as the temperature is lowered to zero. In particular, it was argued [2,8] that the second-order correction to the grand potential can be divergent when the zeroth-order wave function is degenerate, whereas the same correction is finite at zero temperature. These predictions have been confirmed both analytically and

numerically [9,10]. Other instances of the breakdown of diagrammatic perturbation theory have also been reported [11–19].

In this article, we reveal and analyze several additional pathological behaviors of the Feynman-Dyson diagrammatic perturbation expansions of one-particle many-body Green's functions (MBGF) or Feynman propagators [20–29]. Our analysis is based on the *ab initio* electron propagators for molecules [30–71] as we can take advantage of several independent methods that can determine the poles and residues of their exact (finite-basis-set) Green's functions [72] as well as the algorithm that can evaluate perturbation corrections to their frequency-dependent self-energy and Green's function at any arbitrary order and frequency [64]. The conclusions drawn here, however, should be valid for other systems that are studied by the same theory, such as anharmonic molecular vibrations [73], anharmonic lattice vibrations [74], energy bands in solids [75–81], finite nuclei, and nuclear matter [82–84].

^{*}sohirata@illinois.edu

Specifically, we show that (i) the perturbation expansion of the frequency-dependent self-energy is nonconvergent at the exact self-energy in many domains of frequency. (ii) An odd-perturbation-order self-energy has a qualitatively wrong shape except in the central domain that encloses zero frequency and most principal roots. While the diagonal exact self-energy is monotonically decreasing within each frequency bracket separated by its singularities, the diagonal odd-order self-energy can be convex, concave, or monotonically increasing in each bracket. As a result, many roots of the corresponding Dyson equation can be complex and thus nonphysical; when they are real, the corresponding residue can fall outside of the valid range of zero to one. (iii) An even-order self-energy may have a qualitatively correct, monotonically decreasing form within each bracket demarcated by its singularities. However, they become more and more vertical with increasing perturbation order, predicting numerous “phantom” poles with zero residues.

None of these problems is detected in the exact self-energy or Green’s function, which can be determined (within a finite basis set) by the full configuration-interaction (FCI) or, equivalently, by the full equation-of-motion coupled-cluster (EOM-CC) method [64,72], which may be viewed as an infinite resummation of the Feynman-Dyson perturbation corrections [63]. However, (iv) other infinite partial resummations of propagator diagrams by vertex or edge renormalization such as the Tamm-Dancoff approximation [29,33,82,84–87] and self-consistent second-order Green’s function method [8,18,88–98] are shown to exhibit the same pathologies with even greater severity.

We also elucidate (v) the cause of the nonconvergence to be the fact that the definition of the Green’s function is non-analytic at many frequencies, where the radius of convergence of its Taylor expansion is zero. In other words, the breakdown of the Feynman-Dyson perturbation theory occurs due to a purely mathematical reason, i.e., that the rational-function form of the exact Green’s function does not lend itself to a converging Taylor expansion, even though physical information such as the converging perturbation expansions of all state energies is actually contained in it. Consequently, (vi) the general mathematical technique of Padé approximants [34,99–107], which are power-series expansions of a rational function, can extract this physical information, systematically and rapidly restoring the correct shape and poles of the Green’s function.

Nevertheless, the nonconvergence of the Feynman-Dyson perturbation theory, despite being the mathematical foundation of quantum field theory [20–24,27], poses difficulties when applying higher-than-second-order approximations to the Green’s function methods that are predicated on the knowledge of all poles and residues. Such methods include the Galitskii-Migdal identity [108–111], self-consistent Green’s function methods [8,18,88–98], and some models of the algebraic diagrammatic construction (ADC) [51,52,112,113]. The Luttinger-Ward functional [8,15,16,19,88,89,114], which serves as a basis of the self-consistent Green’s function methods and dynamical mean-field theory (DMFT) [115], may also be negatively impacted by these failures.

II. EXACT FEYNMAN PROPAGATOR

A. Formalisms

An electron Feynman propagator is defined in the time (t) domain as a time-ordered sum of Green’s functions,

$$G_{pq}(t) = i\theta(-t)\langle\Psi_0|\hat{p}^\dagger \exp\{i(\hat{H} - E_0)t\} \hat{q}|\Psi_0\rangle - i\theta(t)\langle\Psi_0|\hat{q} \exp\{-i(\hat{H} - E_0)t\} \hat{p}^\dagger|\Psi_0\rangle, \quad (1)$$

where $\theta(t)$ is the Heaviside step function, Ψ_0 and E_0 are the exact wave function and energy for the N -electron ground state, and \hat{p}^\dagger and \hat{q} are the electron creation and annihilation operators. It describes the probability of an electron (hole) traveling from the p th (q th) to q th (p th) spin-orbital in time t ($-t$).

A Fourier transform of Eq. (1) yields the electron propagator in the frequency (ω) domain,

$$G_{pq}(\omega) = \sum_I^{\text{IP}} \frac{\langle\Psi_0|\hat{p}^\dagger|\Psi_I\rangle\langle\Psi_I|\hat{q}|\Psi_0\rangle}{\omega - E_0 + E_I - i\eta} + \sum_A^{\text{EA}} \frac{\langle\Psi_0|\hat{q}|\Psi_A\rangle\langle\Psi_A|\hat{p}^\dagger|\Psi_0\rangle}{\omega - E_A + E_0 + i\eta}, \quad (2)$$

where η is a positive infinitesimal, I sums over all $N - 1$ electron exact states, and A runs over all $N + 1$ electron exact states. (Here, η is just mathematical convenience and has nothing to do with a lifetime of spectral bandwidth, which are infinite and zero, respectively. Nonetheless, the present formulation can be applied to solids without any modification, in which a spectral bandwidth emerges naturally and rigorously as a manifold of closely packed δ -function peaks.)

The first term diverges whenever ω coincides with an exact ionization potential (IP), whereas the second term has a pole at an exact electron-attachment energy (EA), apart from their signs. The primary utility of the electron propagator for molecules and solids is the direct determination of IPs and EAs (or electron-correlated energy bands) for both principal (Koopmans) and satellite (shakeup or non-Koopmans) states. It should be noted, however, the distinction between the principal and satellite roots is not definite, and we use these labels rather loosely in this article.

The exact self-energy $\Sigma(\omega)$ is defined by the Dyson equations,

$$\begin{aligned} G(\omega) &= G^{(0)}(\omega) + G^{(0)}(\omega)\Sigma(\omega)G(\omega) \\ &= G^{(0)}(\omega) + G^{(0)}(\omega)\Sigma(\omega)G^{(0)}(\omega) \\ &\quad + G^{(0)}(\omega)\Sigma(\omega)G^{(0)}(\omega)\Sigma(\omega)G^{(0)}(\omega) + \dots, \end{aligned} \quad (3)$$

with the zeroth-order Green’s function given by

$$G_{pq}^{(0)}(\omega) = \sum_i^{\text{occ.}} \frac{\delta_{pi}\delta_{qi}}{\omega - \epsilon_i - i\eta} + \sum_a^{\text{vir.}} \frac{\delta_{pa}\delta_{qa}}{\omega - \epsilon_a + i\eta}, \quad (5)$$

where “occ.” and “vir.” stand for occupied and virtual spin-orbitals in the N electron ground state of a mean-field theory such as the Hartree-Fock (HF) theory, and ϵ_p denotes the p th canonical spin-orbital energy. Throughout this article, we adhere to the convention [116] that i, j, k , and l label occupied spin-orbitals, a, b, c , and d virtual spin-orbitals, and p and q

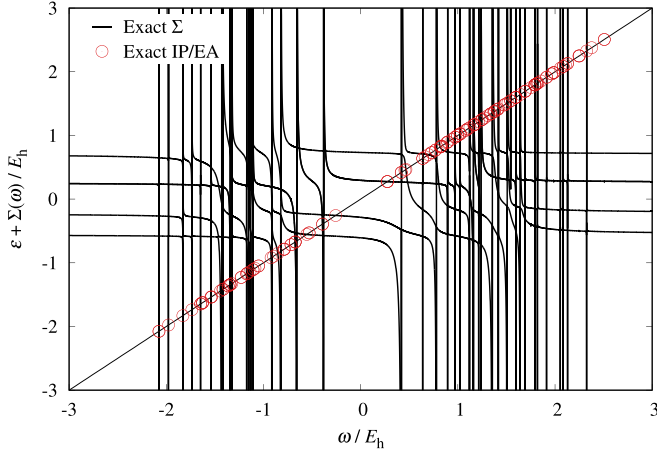


FIG. 3. Eigenvalues of the exact $\epsilon + \Sigma(\omega)$ as a function of ω for the BH molecule. The exact IPs and EAs obtained by FCI are superposed (as open circles), and they coincide with the roots of the inverse Dyson equation, occurring at the intersections of the eigenvalues with the diagonal ω line; see Eq. (9).

lack matching poles, but they correspond either to the poles of other elements of $\mathbf{G}(\omega)$ or to nearly vertical poles (with nearly zero residues) that have fallen through the ω mesh used for plotting Fig. 2. These are merely some minor issues associated with the algorithms adopted, and there is no pathological behavior detectable in the exact MBGF, which reproduces the results of FCI.

Figure 3 plots eigenvalues of the exact $\epsilon + \Sigma(\omega)$ matrix as functions of ω . Of the six eigenvalues the lowest one is not visible in this plot and the second and third highest ones are degenerate. Each root of the inverse Dyson equation is expected at an intersection of the eigenvalues of $\epsilon + \Sigma(\omega)$ and the diagonal line ω as per Eq. (9). In fact, the IPs and EAs obtained from FCI are seen to occur precisely at these intersections. The few IPs and EAs appearing to occur away from any intersection are likely due to the nearly vertical poles with nearly zero residues, which are thus undetected by an ω mesh. This issue may be a weakness of the graphical method [122,123] of solving the inverse Dyson equation if one is concerned with determining all roots, but it by no means signals any fundamental problem in the theory.

Figure 4 is a histogram of the poles of the exact $\mathbf{G}(\omega)$; the height of each impulse is the corresponding residue [Eq. (10)]. There are numerous poles outside of this graph (see Sec. IV). Generally, the poles and residues can be determined by the graphical [122,123] or arrow-matrix diagonalization method [122]. In this study, we used neither; the poles $\{\omega_q\}$ in their entirety were first obtained by FCI [72] and verified by substitution to the inverse Dyson equation [Eq. (9)]. The residue at $\omega = \omega_q$ was then computed by evaluating Eq. (10). The derivative of Σ with respect to ω can be taken analytically by

$$\left(\frac{\partial \Sigma(\omega)}{\partial \omega}\right)_{\omega_q} = \mathbf{1} + \{\mathbf{G}(\omega)\}^{-1} \left(\frac{\partial \mathbf{G}(\omega)}{\partial \omega}\right)_{\omega_q} \{\mathbf{G}(\omega)\}^{-1}, \quad (16)$$

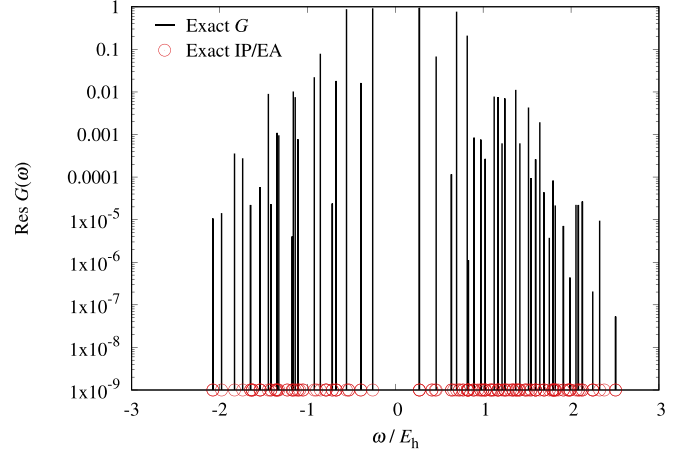


FIG. 4. Residues at poles ω of the exact $\mathbf{G}(\omega)$ for the BH molecule. The exact IPs and EAs obtained by FCI are superposed (as open circles).

with

$$\begin{aligned} \frac{\partial G_{pq}(\omega)}{\partial \omega} = & - \sum_I^{\text{IP}} \frac{\langle \Psi_0 | \hat{p}^\dagger | \Psi_I \rangle \langle \Psi_I | \hat{q} | \Psi_0 \rangle}{(\omega - E_0 + E_I)^2} \\ & - \sum_A^{\text{EA}} \frac{\langle \Psi_0 | \hat{q} | \Psi_A \rangle \langle \Psi_A | \hat{p}^\dagger | \Psi_0 \rangle}{(\omega - E_A + E_0)^2}. \end{aligned} \quad (17)$$

However, the last expression is ill-conditioned at every pole $\omega = \omega_q$. We therefore approximated this derivative as an average of the derivatives at $\omega = \omega_q \pm 10^{-9} E_h$.

The residues thus obtained correctly fall in the range of zero to one for the BH molecule. They also satisfy the particle number sum rule [Eq. (11)] and Galitskii-Migdal identity [Eq. (12)] with the precision of $10^{-7} E_h$ and $10^{-6} E_h$, respectively, which is a numerical manifestation of the fact that the exact Green's function obeys the Baym-Kadanoff conservation laws [29,88,89,93]. Very many tiny contributions from satellite roots are crucial for these identities to be accurately satisfied, and including only principal roots results in severe errors even in this tiny system.

Overall, the exact finite-basis-set Green's function and self-energy are well-behaved, satisfying conservation laws and yielding results that are in exact numerical agreement with alternative methods such as FCI or full EOM-CC. Therefore, the pathological behaviors we are about to discuss are exclusively ascribed to their perturbation expansions.

III. FEYNMAN-DYSON DIAGRAMMATIC PERTURBATION EXPANSION OF PROPAGATOR

A. Formalisms

In most applications, both the Green's function and self-energy are expanded in perturbation series. In this article, perturbation corrections are denoted by symbols prefixed with δ with its order given as the parenthesized superscript:

$$\mathbf{G}(\omega) = \mathbf{G}^{(0)}(\omega) + \delta \mathbf{G}^{(1)}(\omega) + \delta \mathbf{G}^{(2)}(\omega) + \dots, \quad (18)$$

$$\Sigma(\omega) = \delta \Sigma^{(1)}(\omega) + \delta \Sigma^{(2)}(\omega) + \delta \Sigma^{(3)}(\omega) + \dots. \quad (19)$$

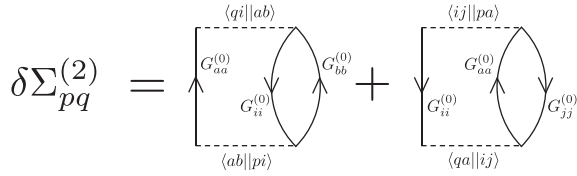


FIG. 5. The second-order self-energy.

Cumulative approximations to \mathbf{G} or Σ through order n are denoted by symbols without a δ prefix.

$$\mathbf{G}^{(n)}(\omega) \equiv \mathbf{G}^{(0)}(\omega) + \sum_{i=1}^n \delta\mathbf{G}^{(i)}(\omega), \quad (20)$$

$$\Sigma^{(n)}(\omega) \equiv \sum_{i=1}^n \delta\Sigma^{(i)}(\omega), \quad (21)$$

In this article, MBGF(n) refers to the n th-order Feynman-Dyson perturbation approximation to one-particle many-body Green's function theory, seeking the poles and residues by solving the inverse Dyson equation with $\Sigma^{(n)}$. On the other hand, the exact MBGF is synonymous to FCI as demonstrated in Sec. II. MBGF(n) is diagrammatically linked [64] and thus size-consistent [124]. Its *ab initio* computer implementations (based on Gaussian-type orbitals [121]) can therefore be applied to both electrons [81] and phonons [74] in solids.

The perturbation corrections are usually stipulated diagrammatically [25–29], but they are also defined [64] algebraically as

$$\delta\mathbf{G}^{(n)}(\omega) = \frac{1}{n!} \left. \frac{\partial^n \mathbf{G}(\omega; \lambda)}{\partial \lambda^n} \right|_{\lambda=0}, \quad (22)$$

$$\delta\Sigma^{(n)}(\omega) = \frac{1}{n!} \left. \frac{\partial^n \Sigma(\omega; \lambda)}{\partial \lambda^n} \right|_{\lambda=0}, \quad (23)$$

$$\begin{aligned} \delta\Sigma_{pq}^{(2)}(\omega) &= (-1)^1 i \sum_i^{\text{occ.}} \sum_{a<b}^{\text{vir.}} \int_{-\infty}^{\infty} \frac{d\omega_a}{2\pi} \int_{-\infty}^{\infty} \frac{d\omega_i}{2\pi} (-i)\langle qi||ab\rangle (-i)\langle ab||pi\rangle iG_{aa}^{(0)}(\omega_a) iG_{ii}^{(0)}(\omega_i) iG_{bb}^{(0)}(\omega + \omega_i - \omega_a) \\ &\quad + (-1)^1 i \sum_a^{\text{vir.}} \sum_{i<j}^{\text{occ.}} \int_{-\infty}^{\infty} \frac{d\omega_i}{2\pi} \int_{-\infty}^{\infty} \frac{d\omega_a}{2\pi} (-i)\langle qa||ij\rangle (-i)\langle ij||pa\rangle iG_{ii}^{(0)}(\omega_i) iG_{aa}^{(0)}(\omega_a) iG_{jj}^{(0)}(\omega + \omega_a - \omega_i) \quad (24) \\ &= \frac{1}{2} \sum_i^{\text{occ.}} \sum_{a,b}^{\text{vir.}} \frac{\langle qi||ab\rangle \langle ab||pi\rangle}{\omega + \epsilon_i - \epsilon_a - \epsilon_b} + \frac{1}{2} \sum_{i,j}^{\text{occ.}} \sum_a^{\text{vir.}} \frac{\langle qa||ij\rangle \langle ij||pa\rangle}{\omega + \epsilon_a - \epsilon_i - \epsilon_j}, \quad (25) \end{aligned}$$

where an occupied spin-orbital index and the imaginary unit both denoted by “ i ” should be distinguished, and $\langle pq||rs\rangle$ is an antisymmetrized two-electron integral [116,121]. Second-order many-body Green's function method [MBGF(2)] solves the inverse Dyson equation [Eq. (9)] with this $\Sigma^{(2)}$. Since the roots of this equation occur at the intersection of $\epsilon + \Sigma^{(2)}(\omega)$ and ω , they can never be divergent even though the method is perturbative.

The second-order correction to the Green's function is then described by the same diagram as the second-order self-energy, but appended with long dangling edges, as shown in Fig. 6. It is important to recognize that the roots of the inverse

where $\mathbf{G}(\omega; \lambda)$ and $\Sigma(\omega; \lambda)$ are the exact (i.e., FCI) values of the respective quantities for a perturbation-scaled Hamiltonian $\hat{H} = \hat{H}^{(0)} + \lambda\hat{V}^{(1)}$. The zeroth-order Hamiltonian $\hat{H}^{(0)}$ corresponds to the $\mathbf{G}^{(0)}$ of Eq. (5). Here, we adopt the HF theory as the zeroth order, which implies $\delta\Sigma_{pq}^{(1)} = 0$ [64].

These λ derivatives can be taken either numerically or analytically. From the former, we obtain benchmark data of the perturbation corrections at several low orders [64]. From the latter, we derive recursions of $\delta\mathbf{G}^{(n)}$ and $\delta\Sigma^{(n)}$ in the style of Rayleigh-Schrödinger perturbation theory, which can then be implemented into a general-order algorithm [64]. This strategy was used in this study. The recursions also justify the diagrammatic rules through the linked-diagram and irreducible-diagram theorems in a time-independent picture [64].

While the time-independent picture is more mathematically transparent and systematically extensible to arbitrarily high orders [64], the time-dependent one may be more appealing to our intuition and expedient [26,27]. In the latter, for example, the second-order self-energy is stipulated diagrammatically as in Fig. 5. It graphically describes the process in which (i) a mean-field particle (hole) scatters another particle out of its mean-field state, thereby creating a hole, at one time; (ii) all three particles and holes propagate in their respective mean-field potentials, i.e., driven by the mean-field propagator $\mathbf{G}^{(0)}$; and (iii) the particle-hole pair recombines at another time. The numerical value of $\delta\Sigma^{(2)}$, which is related to the probability of this overall process, is the product of the probabilities of the constituent scattering and propagation events summed over all possible times and positions of their occurrences. Consulting with Table 4.3 of Mattuck [26], we can then evaluate it as

Dyson equation with $\Sigma^{(2)}$ are *not* the poles of this $\mathbf{G}^{(2)}$; rather, they are the poles of $\mathbf{G}^{\text{Dyson}(2)}$ (using the nomenclature of Holleboom and Snijders [110]) defined by

$$\mathbf{G}^{\text{Dyson}(n)}(\omega) = \{\omega\mathbf{1} - \epsilon - \Sigma^{(n)}(\omega)\}^{-1}, \quad (26)$$

in analogy to Eq. (7) and in accordance with the Dyson equations [Eqs. (3) and (4)]. Therefore, as shown in Fig. 7, $\mathbf{G}^{\text{Dyson}(2)}$ is a *bold-line* Green's function (just like the one appearing in Fig. 1), which includes an infinite-order correction through repeated actions of $\Sigma^{(2)}$. In this sense, MBGF(n) is an infinite-order theory for IPs and EAs even for a finite perturbation order n .

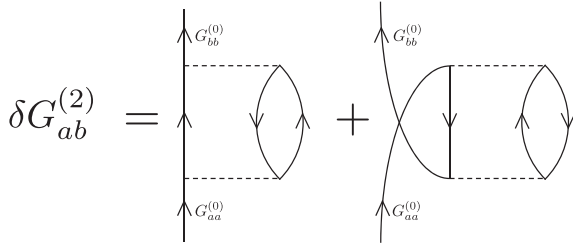


FIG. 6. The second-order Green's function.

Likewise, the third-order self-energy is evaluated from its diagrams [49] as

$$\begin{aligned} \delta\Sigma_{pq}^{(3)}(\omega) &= \frac{1}{4} \sum_i^{\text{occ.}} \sum_{a,b,c,d}^{\text{vir.}} \frac{\langle qi||ab\rangle\langle ab||cd\rangle\langle cd||pi\rangle}{(\omega + \epsilon_i - \epsilon_a - \epsilon_b)(\omega + \epsilon_i - \epsilon_c - \epsilon_d)} \\ &\quad - \frac{1}{4} \sum_{i,j,k,l}^{\text{occ.}} \sum_a^{\text{vir.}} \frac{\langle qa||ij\rangle\langle ij||kl\rangle\langle kl||pa\rangle}{(\omega + \epsilon_a - \epsilon_i - \epsilon_j)(\omega + \epsilon_a - \epsilon_k - \epsilon_l)} \\ &\quad + (16 \text{ terms}), \end{aligned} \quad (27)$$

corresponding to the diagrammatic equation in Fig. 8. It may be noticed that the functional form of $\delta\Sigma^{(3)}$ with respect to ω is different from that of $\delta\Sigma^{(2)}$ [Eq. (25)]; poles in $\delta\Sigma^{(3)}$ are second order, while $\delta\Sigma^{(2)}$ and the exact Green's function have only first-order poles. This difference has a grave consequence on the roots of the inverse Dyson equation, which we now discuss.

B. Numerical results

In Fig. 9 are plotted diagonal elements of the second- and third-order self-energies, $\epsilon_3 + \Sigma_{33}^{(2)}$ and $\epsilon_3 + \Sigma_{33}^{(3)}$, for the third orbital (HOMO) of the BH molecule [64]. Intersections of these functions with the diagonal ω line (also drawn) are the roots of the corresponding inverse Dyson equations in the diagonal approximation. See Eq. (13).

The second-order self-energy has qualitatively the same functional form as the exact self-energy (Fig. 3) in that they are both separated by singularities into consecutive ω brackets, within each of which they are monotonically decreasing and shoulder-like. The singularities of $\Sigma^{(2)}$, i.e., the boundaries of the brackets, occur at the two-particle-one-hole (2p1h) ($\epsilon_a + \epsilon_b - \epsilon_i$) and two-hole-one-particle (2h1p) ($\epsilon_i + \epsilon_j - \epsilon_a$)

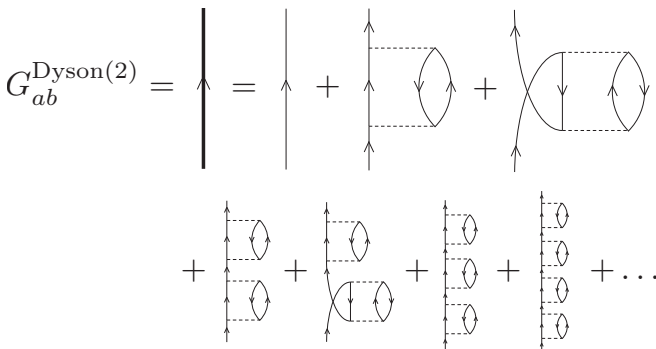


FIG. 7. The second-order bold-line Green's function.

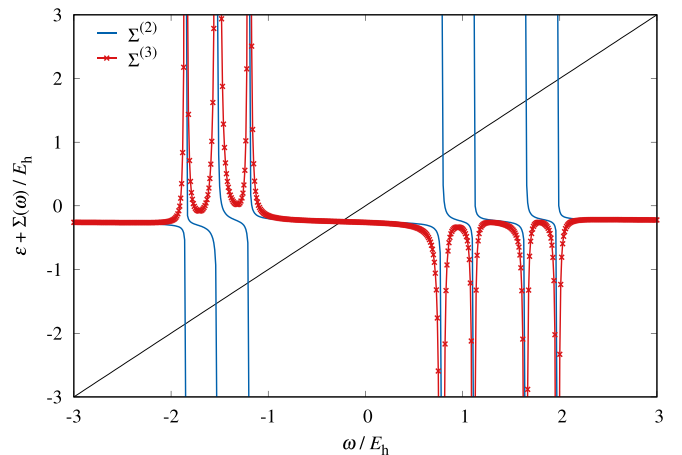
$$\delta\Sigma_{pq}^{(3)} = \text{[Diagram 1]} + \text{[Diagram 2]} + (16 \text{ diagrams})$$

FIG. 8. The third-order self-energy. See Appendix 1 of Öhrn and Born [49] for a complete list.

HF orbital energy differences according to Eq. (25). In each bracket, a diagonal element of the second-order self-energy intersects the diagonal ω line exactly once, just as the exact self-energy does [122]. The second-order inverse Dyson equation has the principal root for the HOMO at around $-0.25E_h$ in the central bracket enclosing $\omega = 0$, where the self-energy is relatively flat and whose residue is close to unity [see Eq. (14)]. The self-energy also intersects the ω line in each of the other brackets, typically on a near-vertical part of its shoulder-like curve. This means that these satellite roots coincide with the 2p1h or 2h1p HF orbital energy differences, accounting for little to no electron-correlation effects, and have near zero residues. Nevertheless, MBGF(2) is overall well-behaved.

The third-order self-energy, in contrast, has a qualitatively wrong functional form. It is still separated by the same 2p1h and 2h1p singularities into the identical set of brackets [see Eq. (27)], but within each bracket, $\Sigma^{(3)}$ is either concave or convex except in the central bracket, where it is correctly monotonically decreasing. Consequently, in the domain of ω of Fig. 9, the third-order inverse Dyson equation has only one real root, i.e., the principal root for the HOMO at around $-0.25E_h$. The different functional forms of the self-energy between the second and third orders can be easily rationalized by their algebraic definitions, Eqs. (25) and (27). The second-order self-energy (as well as the exact self-energy) has only first-order poles, whereas the third-order self-energy features up to second-order poles.

Figure 10 plots the second- and third-order bold-line Green's functions. The $G^{\text{Dyson}(2)}$ has the same overall appearance as the exact Green's function (Fig. 2). It exhibits a "fat" pole at the principal root for the HOMO and several "thin" poles at satellite roots. In contrast, $G^{\text{Dyson}(3)}$ displays

FIG. 9. The third diagonal elements of $\epsilon + \Sigma^{(2)}(\omega)$ and $\epsilon + \Sigma^{(3)}(\omega)$ as functions of ω for the BH molecule.

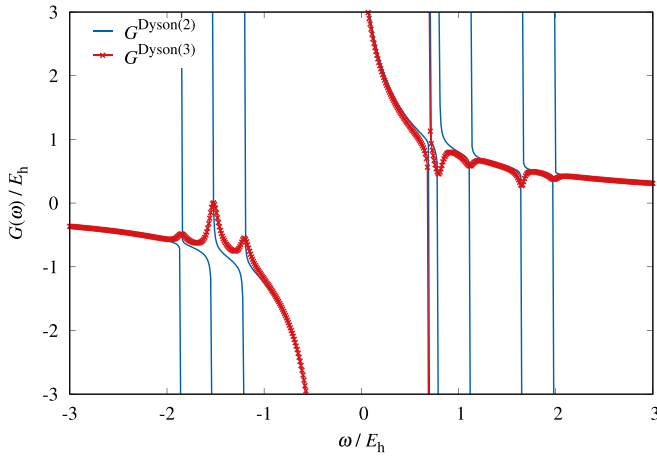


FIG. 10. The third diagonal elements of $G^{\text{Dyson}(2)}(\omega)$ and $G^{\text{Dyson}(3)}(\omega)$ as functions of ω for the BH molecule.

only undulations but no poles at the frequencies where satellite roots are expected. The diagonal element of $G^{\text{Dyson}(3)}$ for the HOMO has poles only at the principal roots at around $-0.25E_h$ and $0.69E_h$, which is consistent with Fig. 9. (Although there is no root at around $0.69E_h$ in MBGF(3) according to Fig. 9, this principal EA root has a considerable mixing with other roots with its residue being only 0.77, which is why it creates a pole in $G_{33}^{\text{Dyson}(3)}$.) Hence, the perturbative Green's functions confirm the absence of many real satellite roots at MBGF(3).

Where have the missing satellite roots of MBGF(3) gone? We believe that they have gone to the complex space, although no attempt has been made to determine them numerically.

Figure 11 plots all six diagonal elements of $\epsilon + \Sigma^{(3)}$ in a full domain of ω . For the HF occupied orbitals 1, 2, 3, and virtual orbital 6 (labeled in the increasing order of energy), which transform as a_1 in the C_{4v} subgroup, there are twelve (12) 2p1h and 2h1p singularities or thirteen (13) brackets. Hence, the second- and third-order inverse Dyson equations in the diagonal approximation are a thirteenth-order polynomial equation for each of these a_1 orbitals. The second-order equation has one principal root and twelve satellite roots (13 in total), all of which are real.

On the other hand, as Fig. 11 indicates, the diagonal element of the third-order self-energy for orbital 1 intersects the ω line only nine times, meaning that its diagonal third-order inverse Dyson equation has at least nine real roots (some may be degenerate) out of 13 roots in total. The remaining four (or less in the case of degeneracy) roots must be complex. For orbitals 2, 3, and 6, the number of intersections is three and hence the third-order equation for each of these orbitals has three (or more) real and ten (or fewer) complex roots.

The HF virtual orbitals 4 and 5 transform as e of C_{4v} and are degenerate. There are nine 2p1h and 2h1p singularities or ten brackets for these orbitals. The diagonal second- and third-order inverse Dyson equations are therefore a tenth-order polynomial equation. The second-order equation has ten real

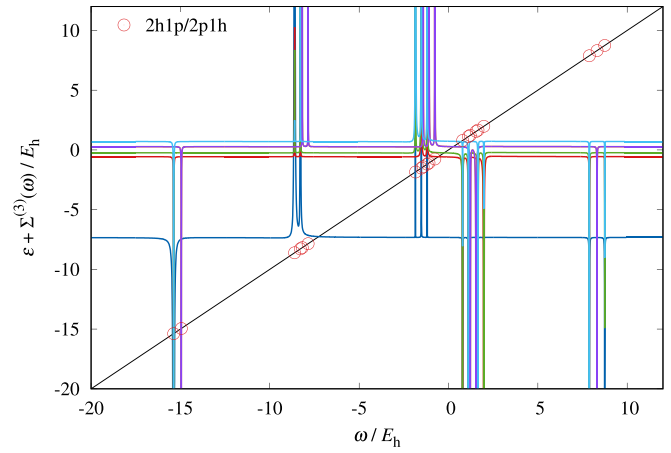


FIG. 11. All six diagonal elements of $\epsilon + \Sigma^{(3)}(\omega)$ as functions of ω for the BH molecule. The fourth and fifth elements are degenerate, appearing as one curve. The 2p1h and 2h1p HF orbital energy differences are superposed.

roots, but as seen in Fig. 11, the third-order self-energy intersects the ω line only three times, implying that the third-order equation has four or more real roots and six or less complex roots (since complex roots of a polynomial equation with real coefficients occur in complex-conjugate pairs, at least one of the real roots must be degenerate).

From Fig. 11, it can furthermore be observed that real satellite roots of MBGF(3) tend to appear as *nearly* degenerate pairs or doublets (which is distinct from the exact degeneracy of roots mentioned in the foregoing paragraphs). One of a doublet corresponds to an intersection of a sharply falling part of the self-energy with the ω line and thus has a positive infinitesimal residue, while the other intersection occurs at an almost vertically *rising* part of the self-energy, whose residue is negative and thus nonphysical. These intersections almost coincide with bare 2p1h or 2h1p HF orbital energy differences, thus accounting for little to no correlation effects.

These observations can be generalized to all orders. One can expect an even-order self-energy to have the same qualitatively correct functional form as the exact self-energy, while the third and higher odd-order self-energies to display qualitatively wrong functional forms. This is borne out in Fig. 12, in which the eighth- and ninth-order self-energies are plotted along with the $(n+1)hnp$ and $(n+1)pnh$ HF orbital energy differences. In Fig. 13, all six diagonal elements of $\Sigma^{(8)}$ are drawn in a full domain of ω .

In Fig. 12, the odd-order $\Sigma^{(9)}$ has a concave or convex shape in each bracket separated by its singularities and, as a result, intersects the ω line only once in the shown ω domain. In the same figure, the exact self-energy and exact IPs and EAs are overlaid, indicating that there should be far more intersections than just one in this ω domain. The missing satellite roots of MBGF(9) are believed to be complex.

The eighth-order self-energy is less problematic in this regard, but it comes with new issues. Its brackets are separated by the 2p1h, 2h1p, 3p2h, 3h2p, 4p3h, 4h3p, 5p4h, and 5h4p HF orbital energy differences. The even-order $\Sigma^{(8)}$

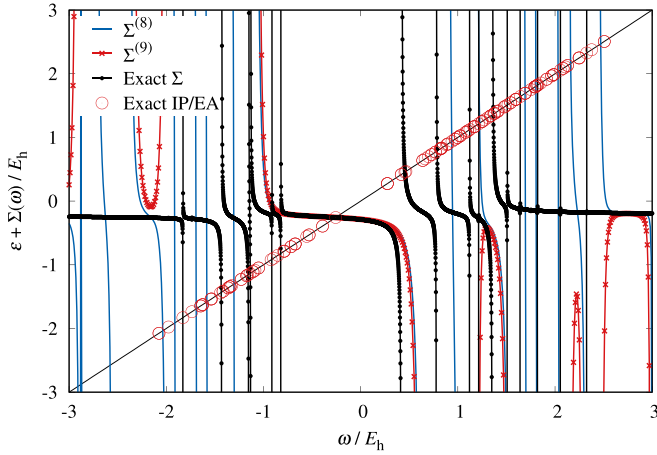


FIG. 12. The third diagonal elements of the exact $\epsilon + \Sigma(\omega)$ and $\epsilon + \Sigma^{(n)}(\omega)$ ($n = 8, 9$) as functions of ω for the BH molecule. The exact IPs and EAs obtained by FCI are superposed.

is technically monotonically decreasing within each bracket, but it decreases so sharply in many brackets that it consists of near vertical lines, where it intersects the ω line, as seen in Figs. 12 and 13. Therefore, although MBGF(8) has real roots, most of them are phantom poles with zero residues, occurring somewhere near the midpoint of each bracket.

Consequently, the distribution of the roots of the eighth-order inverse Dyson equation (the intersections of $\epsilon + \Sigma^{(8)}$ with the ω line in Fig. 13) matches more closely the mere HF orbital energy differences (the green diamond plots in the same figure) than the exact roots (the red circles). Even though these MBGF(8) roots are not equal to the HF orbital energy differences, they are confined within brackets demarcated by these differences, which form dense manifolds, and tend to agree with them. These satellite roots therefore include little to no electron-correlation effects, not to mention that most of them hardly exist physically because their residues are zero.

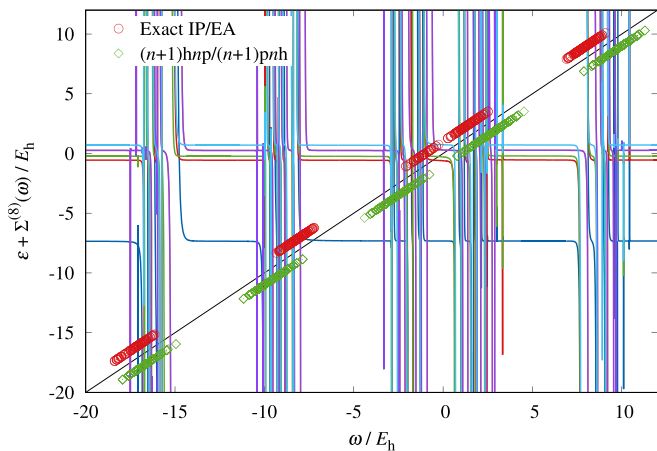


FIG. 13. All six diagonal elements of $\epsilon + \Sigma^{(8)}(\omega)$ as functions of ω for the BH molecule. The $(n+1)pnh$ and $(n+1)hnp$ HF orbital energy differences ($n = 1, 2, 3$) as well as the exact IPs and EAs obtained by FCI are superposed.

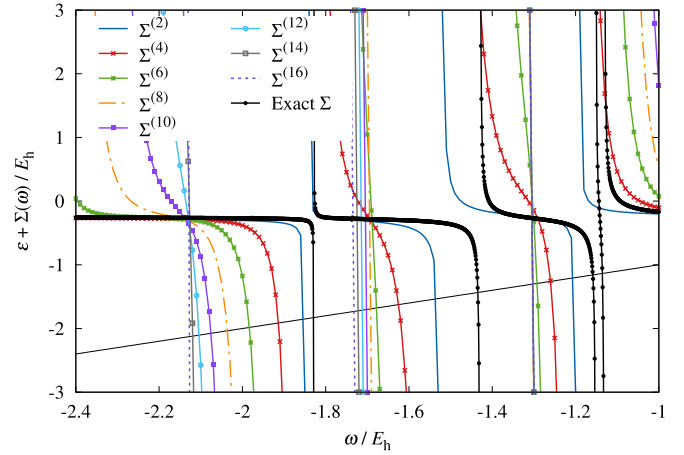


FIG. 14. The third diagonal elements of the exact $\epsilon + \Sigma(\omega)$ and $\epsilon + \Sigma^{(n)}(\omega)$ ($n = 2, 4, 6, \dots, 16$) as functions of ω for the BH molecule.

These are disheartening results considering that the eighth- or ninth-order perturbation approximations are usually numerically exact in most other physics cases [105, 107].

Our numerical tests for other molecules (not shown) have indicated that an odd-order self-energy can generally be concave, convex, or even monotonically *increasing* within a bracket. Therefore, an odd-order inverse Dyson equation can have real roots (sometimes more than one in a bracket), but the corresponding residues may fall outside the valid range of zero to one.

One might still wonder if selectively raising only the even perturbation order may avoid these pathologies. Figure 14 shows the evolution of $\Sigma^{(2n)}$ as a function of the even perturbation order $2n$. With increasing order, the even self-energy becomes more and more vertical and less shoulder-like with the intersections with the ω line moving away from the correct locations. The corresponding residues will approach zero. Ironically, it appears as though the perturbative self-energies converged at the exact one as the perturbation order is lowered! This figure is the most compelling piece of evidence demonstrating the nonconvergence of the Feynman-Dyson perturbation expansion of the self-energy.

This conclusion does not undermine the utility or rapid convergence of MBGF(n) for roots falling well within the central bracket bounded by the highest 2h1p and lowest 2p1h energies as well as in some other ω domains (see Sec. V A). This bracket typically encloses most principal IPs and EAs (but not core IPs) as well as low-energy satellite poles. Figure 15 is a close-up of the self-energy-versus- ω plot for MBGF(n) ($2 \leq n \leq 11$) as well as the exact MBGF. It shows that the intersections of the self-energy and ω line systematically approach the exact IP for the HOMO.

IV. INFINITE PARTIAL RESUMMATIONS OF PROPAGATOR DIAGRAMS

A. Vertex renormalization

To avoid the malaise of truncated perturbation approximations, infinite partial resummations of diagrams have often

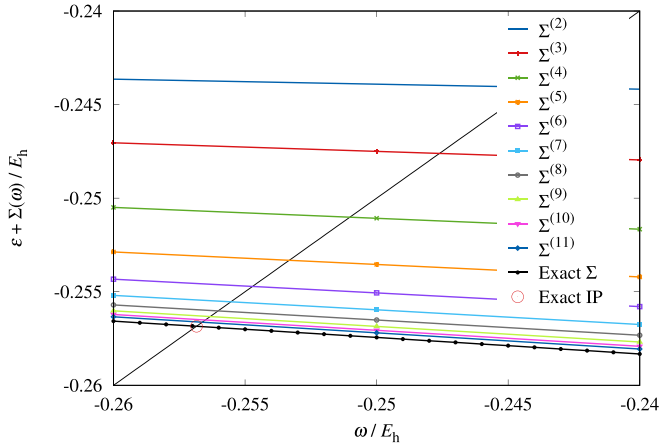


FIG. 15. The third eigenvalues of the exact $\epsilon + \Sigma(\omega)$ and $\epsilon + \Sigma^{(n)}(\omega)$ ($n = 2, 3, 4, \dots, 11$) as functions of ω for the BH molecule. The corresponding exact IP obtained from FCI is superposed.

been invoked. One way of doing this is by renormalizing vertexes.

Here, a “vertex” refers to an instantaneous Coulomb interaction denoted by a horizontal (dashed) line in a Hugenholtz-Shavitt-Bartlett diagram and is equal to an antisymmetrized two-electron integral containing both the direct *and* exchange contributions [116,121]. It should be distinguished from a space-time point of contact between propagator lines and an interaction line in a Goldstone diagram, each standing for only the direct *or* exchange contribution, often used in the *GW* approximations [31,55,58,60,125,126].

In the two-particle-hole Tamm-Dancoff approximation (TDA) [33,85–87], also known as the Brueckner-Hartree-Fock method [82,84] or the T approximation [29], the diagrams of the types in Fig. 16 are summed over up to an infinite order. In this figure, the first two diagrams in the third and fourth lines are “ring” diagrams, and the subsequent two diagrams are “ladder” diagrams. The effect of this infinite resummation can be folded into the bold-line vertexes as appearing in the first line of Fig. 16, which must, in turn, satisfy the diagrammatic equations drawn in Fig. 17. They

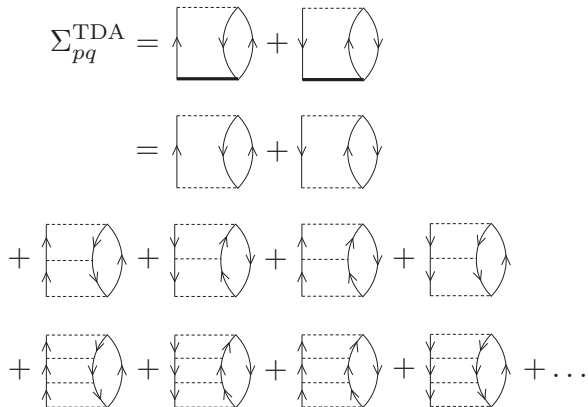


FIG. 16. The self-energy in the TDA approximation.

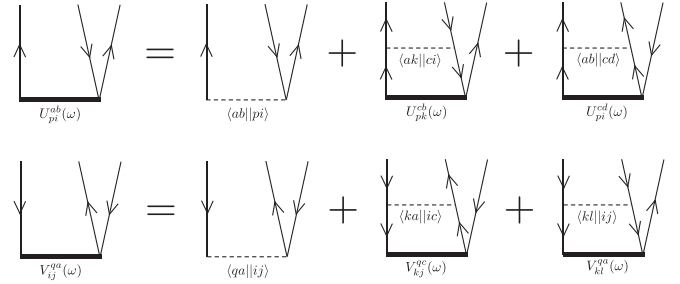


FIG. 17. The diagrammatic equations for the bold-line (renormalized) vertexes of TDA.

take the form of the amplitude equations of coupled-cluster theory [116,127] and are written algebraically as

$$\begin{aligned} (\omega + \epsilon_i - \epsilon_a - \epsilon_b)U_{pi}^{ab}(\omega) &= \langle ab||pi \rangle - P(ab) \sum_{c,k} \langle ak||ci \rangle U_{pk}^{cb}(\omega) \\ &+ \frac{1}{2} \sum_{c,d} \langle ab||cd \rangle U_{pi}^{cd}(\omega), \end{aligned} \quad (28)$$

$$\begin{aligned} (\epsilon_i + \epsilon_j - \omega - \epsilon_a)V_{ij}^{qa}(\omega) &= \langle qa||ij \rangle - P(ij) \sum_{c,k} \langle ka||ic \rangle V_{kj}^{qc}(\omega) \\ &+ \frac{1}{2} \sum_{k,l} \langle kl||ij \rangle V_{kl}^{qa}(\omega), \end{aligned} \quad (29)$$

where $P(ab)$ is an antisymmetrizer [116]. Unlike coupled-cluster theory, where there is only one type of cluster excitation amplitudes (denoted by T), there are two (2p1h and 2h1p) types of modified vertexes whose numerical values are stored in U and V . They represent electron-electron repulsion tempered by screening and other higher-order electron-correlation effects.

Equations (28) and (29) are a system of linear equations, which can, therefore, be ill-conditioned at some ω . In this work, they are solved in an iterative algorithm; i.e., starting with initial guesses of U and V , we substitute them in the right-hand sides of these equations to update U and V in the left-hand sides, and repeat this process until convergence. Therefore, in practice, the highest order of the ring and ladder diagrams that are actually included in the calculation is capped by the number of cycles taken in this iterative solution. Upon convergence, the self-energy is obtained as

$$\Sigma_{pq}^{TDA}(\omega) = \frac{1}{2} \sum_{i,a,b} \langle qi||ab \rangle U_{pi}^{ab}(\omega) - \frac{1}{2} \sum_{i,j,a} \langle ij||pa \rangle V_{ij}^{qa}(\omega). \quad (30)$$

See Ref. [71] for a more efficient and stable algorithm, which, however, is not exempt from the pathological behaviors discussed below, which are deeply rooted in the formalism.

For a total energy, an infinite partial resummation of the corresponding ring and ladder diagrams defines an instance of coupled-cluster theory [116,127] known as linearized coupled-cluster doubles (LCCD) or by other names [128–134]. This method is ω -independent and is thus free

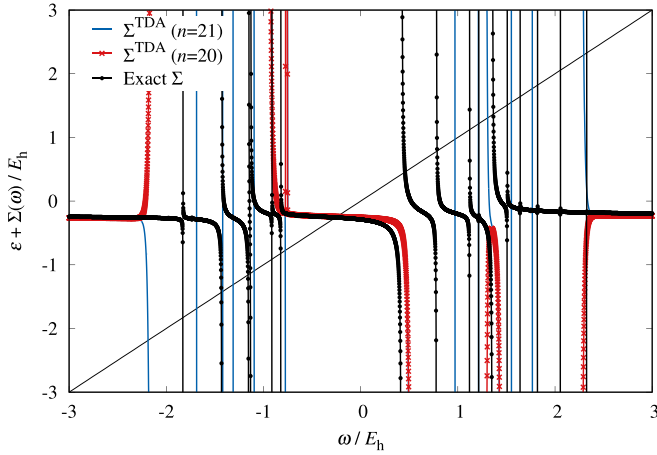


FIG. 18. The exact $\epsilon_3 + \Sigma_{33}(\omega)$ and $\epsilon_3 + \Sigma_{33}^{\text{TDA}}(\omega)$ as functions of ω for the BH molecule. The self-energy in TDA is determined after n cycles of the iterative solution of the amplitude equation, summing over the ring and ladder diagrams through the order $n + 1$.

from the nonconvergence problems discussed here. It instead displays other problems concerning accuracy [135].

Figure 18 compares the self-energy of TDA with the exact self-energy. The former was obtained after either 20 or 21 cycles of the iterative solution of the amplitude equations, summing over ring and ladder diagrams through the order 21 or 22, respectively. As may be expected from the discussion on perturbative self-energies, TDA does not improve the overall appearance of the self-energy; rather, it degrades it.

For instance, Σ^{TDA} obtained after 20 cycles has qualitatively wrong (concave and convex) functional forms (outside the central bracket) because computationally it is a high-odd-order perturbation theory. In contrast, Σ^{TDA} after 21 cycles has qualitatively correct functional forms (as it is a high-even-order perturbation theory), but it consists of numerous vertical lines. Therefore, TDA predicts vastly different IPs and EAs depending on the number of cycles taken in the iterative solution—an artifact of calculations—and is therefore methodologically ill-defined insofar as all roots are sought. Furthermore, both of these conflicting predictions are equally meaningless. After 20 cycles, there are no real roots outside the central bracket in this graph; after 21 cycles, most roots are phantom poles with zero residues. However, this observation should not be misconstrued to mean that TDA is useless; for the principal IPs and EAs, TDA is sound and useful.

It is unsurprising for an infinite partial resummation to inherit the pathologies of higher-order perturbation theory, but this does not imply that the same pathologies plague all vertex renormalization techniques. The equation-of-motion (EOM) formalism of Green’s function theory [37,45–47,83] also involves an infinite partial resummation of diagrams, but it can be recast into a matrix diagonalization [48] and may thus be free from the type of the nonconvergence problems discussed above. The random-phase approximation (RPA) [91], which is a lowest-order member of the EOM approximation series, as well as the related GW approximations [31,55,58,60,125,126] and the frequency-dependent part of the ADC self-energy [51,52], may also be more robust than TDA, although this has not been numerically confirmed nor is the convergence

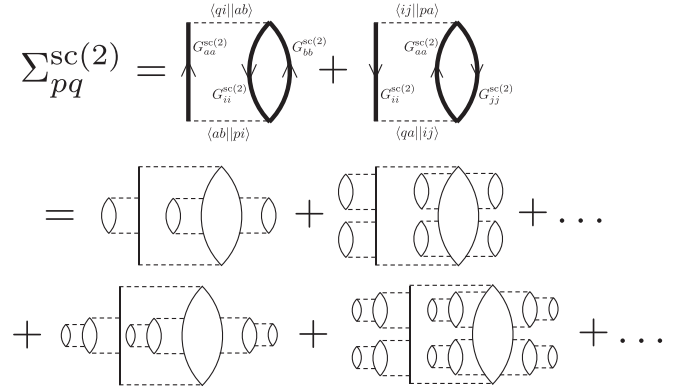


FIG. 19. The self-consistent second-order self-energy.

toward exactness implied. The Parquet method [136] sums over a related, but distinct class of diagrams and hence the present conclusion may not apply, either.

The EOM-CC method [72,137–146] has been argued to be a coupled-cluster Green’s function [147–157] and can also be viewed as an infinite resummation of the propagator diagrams [63]. However, this method does not suffer from the pathologies discussed above; it instead yields IPs and EAs for all (principal, satellite, and even core-ionized [158]) states that are systematically convergent at the exact values with increasing excitation level. EOM-CC separates the IP and EA sectors and its roots are obtained as eigenvalues of a non-Hermitian matrix. Like MBGF(n), the EOM-CC roots are not guaranteed to be real, but they are almost never complex in practice.

It may be inferred that the EOM formalisms of Green’s functions, be they based on perturbation theory or coupled-cluster theory, are more robust and potentially converging because their IPs and EAs are obtained directly as eigenvalues of some matrix rather than by solving an inverse Dyson equation. In other words, the EOM formalisms may attain stability and convergence by divorcing from Green’s function theory in the Feynman-Dyson style. That the exact MBGF and MBGF(2), the latter being the only Feynman-Dyson perturbation order largely exempt from the pathologies, can also be recast into a matrix eigenvalue problem [122] is consistent with this inference.

B. Edge renormalization

Another way of performing a diagram resummation is by renormalizing edges. By replacing all three edges in each diagram of $\delta\Sigma^{(2)}$ by the corresponding bold-line Green’s functions $\mathbf{G}^{\text{Dyson}(2)}$ of Fig. 7, we include an infinite number of “row-house” diagrams appearing in the second line of Fig. 19. Furthermore, if the bold-line Green’s function (designated by $\mathbf{G}^{\text{sc}(2)}$) comes from this very edge-modified self-energy $\Sigma^{\text{sc}(2)}$, as in Fig. 20, we account for another infinite set of “tower” diagrams shown in the third line of Fig. 19. This self-consistency between the self-energy and Green’s function was emphasized by Baym and Kadanoff [29,88,89,93] as an essential ingredient of an approximate MBGF method that obeys conservation laws. (The first-order self-consistent MBGF method is identified as the HF theory [91].) In

chemistry, approximations inspired by this idea are enjoying a revival for its possible ability to describe strong correlation more accurately [96,98,110,159].

In the diagonal approximation, a *self-consistent* second-order self-energy [90] is thus defined by the same equation as Eq. (24) but with each $\mathbf{G}^{(0)}$ replaced by $\mathbf{G}^{\text{sc}(2)}$,

$$\begin{aligned} \Sigma_{pq}^{\text{sc}(2)}(\omega) &= (-1)^1 i \sum_i^{\text{occ.}} \sum_{a < b}^{\text{vir.}} \int_{-\infty}^{\infty} \frac{d\omega_a}{2\pi} \int_{-\infty}^{\infty} \frac{d\omega_i}{2\pi} (-i)\langle qi||ab\rangle (-i)\langle ab||pi\rangle iG_{aa}^{\text{sc}(2)}(\omega_a) iG_{ii}^{\text{sc}(2)}(\omega_i) iG_{bb}^{\text{sc}(2)}(\omega + \omega_i - \omega_a) \\ &+ (-1)^1 i \sum_a^{\text{vir.}} \sum_{i < j}^{\text{occ.}} \int_{-\infty}^{\infty} \frac{d\omega_i}{2\pi} \int_{-\infty}^{\infty} \frac{d\omega_a}{2\pi} (-i)\langle qa||ij\rangle (-i)\langle ij||pa\rangle iG_{ii}^{\text{sc}(2)}(\omega_i) iG_{aa}^{\text{sc}(2)}(\omega_a) iG_{jj}^{\text{sc}(2)}(\omega + \omega_a - \omega_i) \quad (31) \\ &= \frac{1}{2} \sum_i^{\text{occ.}} \sum_{a,b}^{\text{vir.}} \sum_{I(i)}^{\text{IP}} \sum_{A(a),B(b)}^{\text{EA}} \frac{\langle qi||ab\rangle \langle ab||pi\rangle}{\omega + \omega_{I(i)} - \omega_{A(a)} - \omega_{B(b)}} F(\omega_{I(i)}) F(\omega_{A(a)}) F(\omega_{B(b)}) \\ &+ \frac{1}{2} \sum_a^{\text{vir.}} \sum_{i,j}^{\text{occ.}} \sum_{I(i),J(j)}^{\text{IP}} \sum_{A(a)}^{\text{EA}} \frac{\langle qa||ij\rangle \langle ij||pa\rangle}{\omega + \omega_{A(a)} - \omega_{I(i)} - \omega_{J(j)}} F(\omega_{A(a)}) F(\omega_{I(i)}) F(\omega_{J(j)}), \quad (32) \end{aligned}$$

where the occupied spin-orbital index i and the imaginary unit i are to be distinguished, and $\omega_{I(q)} (< 0)$ and $\omega_{A(q)} (> 0)$ are an IP and EA root, respectively, of the inverse Dyson equation in the diagonal approximation, which satisfy

$$\epsilon_q + \Sigma_{qq}^{\text{sc}(2)}(\omega_{I(q)}) = \omega_{I(q)}, \quad (33)$$

$$\epsilon_q + \Sigma_{qq}^{\text{sc}(2)}(\omega_{A(q)}) = \omega_{A(q)}. \quad (34)$$

This *Ansatz* may differ somewhat from those of Van Neck *et al.* [90] or of Dahlen and van Leeuwen [93] because it involves some additional approximations, but these differences have no impact on the following analysis. The corresponding residues are given by

$$F(\omega_{I(q)}) \equiv \text{Res}_{\omega_{I(q)}} G_{qq}^{\text{sc}(2)}(\omega) = \left\{ 1 - \left. \frac{\partial \Sigma_{qq}^{\text{sc}(2)}(\omega)}{\partial \omega} \right|_{\omega_{I(q)}} \right\}^{-1}. \quad (35)$$

These conditions imply the self-consistency,

$$\mathbf{G}^{\text{sc}(2)}(\omega) = \{\omega \mathbf{1} - \epsilon - \Sigma^{\text{sc}(2)}(\omega)\}^{-1}, \quad (36)$$

in the diagonal approximation, although an explicit evaluation of $\mathbf{G}^{\text{sc}(2)}$ is never needed.

In practice, Eqs. (32)–(35) are solved iteratively. In cycle zero ($n = 0$), we use $\mathbf{G}^{(0)}$ in the right-hand side of Eq. (32) and obtain $\Sigma^{\text{sc}(2)}(n = 0) = \Sigma^{(2)}$ in the left-hand side. In cycle one, therefore, the new Green's function is $\mathbf{G}^{\text{Dyson}(2)}$. By determining all of its poles and residues and substituting them back into Eq. (32), we obtain $\Sigma^{\text{sc}(2)}(n = 1)$. In cycle two,

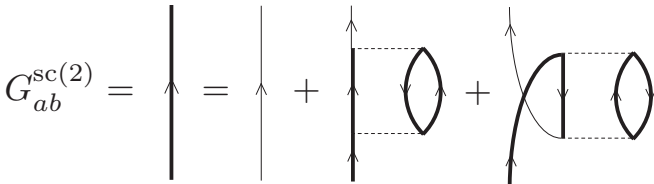


FIG. 20. The diagrammatic equation for the bold-line (renormalized) edge of the self-consistent second-order Green's function method.

similarly, we get $\Sigma^{\text{sc}(2)}(n = 2)$, and so on. In each cycle, figuratively speaking, a “new floor” is added to each of the “tower” diagrams (see the third line of Fig. 19). This calculation quickly becomes intractable because the number of poles increases factorially with iterative cycles. Dahlen and van Leeuwen [93] and Zgid and coworkers [96,97,160,161] devised imaginary-time-dependent (or finite-temperature) algorithms for this method, which seem to overcome this computational intractability.

Figure 21 compares the self-energies of the self-consistent second-order Green's function method after zeroth, first, and second iterative cycles. In each case, the self-energy has the qualitatively correct functional form, i.e., monotonically decreasing in each bracket separated by its singularities. This is because the self-energy expression [Eq. (32)] is isomorphic to the second-order self-energy, which has only the first-order poles. Furthermore, unlike higher-order perturba-

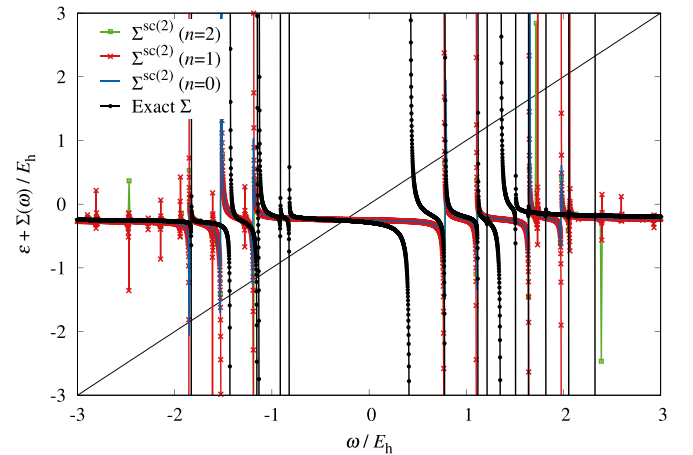


FIG. 21. The exact $\epsilon_3 + \Sigma_{33}(\omega)$ and $\epsilon_3 + \Sigma_{33}^{\text{sc}(2)}(\omega)$ as a function of ω for the BH molecule. The self-consistent self-energy is determined after n cycles of the iterative i th (a th) edge replacement by diagonal bold-line $G_{ii}^{\text{sc}(2)}$ ($G_{aa}^{\text{sc}(2)}$). $\Sigma^{\text{sc}(2)}(n = 0)$ corresponds to the unmodified $\Sigma^{(2)}(\omega)$.

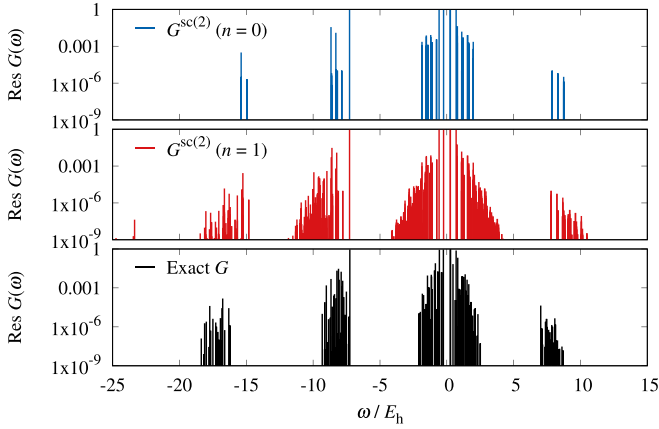


FIG. 22. Residues at poles ω of $G^{\text{sc}(2)}(\omega)$ after n cycles of the self-consistent iteration as well as of the exact $G(\omega)$ for the BH molecule. $G^{\text{sc}(2)}(\omega)$ at $n = 0$ corresponds to $G^{\text{Dyson}(2)}(\omega)$.

tive self-energies or TDA, the brackets' boundaries shift from one cycle to the next, and hence the satellite roots are no longer confined by brackets of the mere HF orbital energy differences.

Nonetheless, this does not mean that these satellite roots are improved. To the contrary, they seem to deteriorate with increasing iterative cycles. Figure 21 shows that in the first and second cycles, new singularities of the self-energy emerge, e.g., in the domain $-3E_h \leq \omega \leq -2E_h$, where there are no corresponding singularities of the exact self-energy. The mechanism by which these spurious singularities multiply rapidly with increasing self-consistent cycles is as follows: In each cycle, the poles of the Green's function define new ω brackets, in each of which there is one root of the inverse Dyson equation. These brackets are demarcated by singularities of the self-energy, Eq. (32), which are the $2p1h$ ($\omega_{A(a)} + \omega_{B(b)} - \omega_{I(i)}$) or $2h1p$ ($\omega_{I(i)} + \omega_{J(j)} - \omega_{A(a)}$) energy differences of the poles of the Green's function (which are no longer the HF orbital energy differences after the first cycle). As the number of poles increases, the number of brackets and thus the number of roots increase extremely rapidly, quickly exceeding the correct total number of roots in a finite basis set. The growth is factorial of the iterative cycle.

Figure 22 shows the poles and residues in the zeroth and first cycles of the self-consistent second-order Green's function method in comparison with the exact poles and residues. The bottom panel is the same as Fig. 4, but shown in a full domain of ω , and the top panel is equivalent to MBGF(2) in the diagonal approximation. In the zeroth cycle ($n = 0$), the distribution of the poles agree reasonably well with that of the exact poles. The number of poles is 72 as compared with 600 exact poles including ones with zero residues. In the first cycle ($n = 1$), the number of poles already reaches 4314, far exceeding the total number (600) of ionized and electron-attached states of a FCI calculation. Furthermore, these spurious poles are not necessarily phantom mathematical roots with zero residues; they have nonzero residues and encroach on the regions where there are no exact poles. In the second cycle ($n = 2$), the number of roots reaches such an astronomical value that our computer code can no longer

handle, and we judged that it was not worthwhile to pursue full self-consistency. This difficulty was recognized by earlier workers, many of whom then decided to abandon this class of methods.

The bold-line self-energy diagrams in Fig. 19 is obtained by cutting one bold line (and then trimming the resulting two dangling lines) [64] of a skeleton diagram of the Luttinger-Ward functional [8,88,89,114] in the zero-temperature limit. Several mathematical difficulties associated with this functional have been reported recently [12,14–19].

V. ANALYSIS

A. Cause of nonconvergence

What is the cause of the nonconvergence? This can be answered by analyzing a model Green's function of the form [9]

$$g(\omega) = \frac{1}{\omega - E_1} + \frac{1}{\omega - E_2} + \frac{1}{\omega - E_3} + \frac{1}{\omega - E_4}, \quad (37)$$

which consists of four poles. These poles are stipulated to be quadratic functions of the perturbation strength λ :

$$E_1 = 1.9 + 0.2\lambda + 0.2\lambda^2, \quad (38)$$

$$E_2 = 0.75 + 0.1\lambda + 0.1\lambda^2, \quad (39)$$

$$E_3 = -1.1 - 0.1\lambda - 0.1\lambda^2, \quad (40)$$

$$E_4 = -2.2 - 0.15\lambda - 0.15\lambda^2. \quad (41)$$

The functional forms of $g(\omega)$ and the E s are arbitrary, but a different choice would lead to essentially the same conclusion.

One can expand $g(\omega)$ in a Taylor series in λ ,

$$g(\omega) = g^{(0)}(\omega) + \lambda g^{(1)}(\omega) + \lambda^2 g^{(2)}(\omega) + \lambda^3 g^{(3)}(\omega) + \dots \quad (42)$$

A truncation of this series after a finite number of terms captures the key features of the Feynman-Dyson perturbation approximations of a Green's function or self-energy.

Figure 23 shows the zeroth-, first-, and second-order Taylor expansions of $g(\omega)$. The exact $g(\omega)$ and its zeroth-order approximation $g^{(0)}(\omega)$ have essentially the same functional forms in that they are separated into consecutive ω brackets by their singularities and within each bracket they are monotonically decreasing functions. The first-order approximation $g^{(0)} + g^{(1)}$ has a qualitatively different functional form, which is convex or concave except in the central bracket. The $g^{(0)} + g^{(1)} + g^{(2)}$ largely restores the same functional form as the exact $g(\omega)$, although the former is a rather poor approximation at many frequencies despite the fact that the second-order Taylor expansions of E_1 through E_4 are exact. These are consistent with the overall patterns of behaviors of *ab initio* odd- and even-order perturbative self-energies observed in the foregoing sections.

Figure 24 extends this analysis to the nineteenth-order Taylor expansion. It is convergent at the exact $g(\omega)$ in some domains of ω , but nonconvergent in the other domains. Generally, convergence is attained within the overlap of the bracket demarcated by the singularities of $g(\omega)$ and the one delineated by the singularities of $g^{(0)}(\omega)$. (The latter singularities are

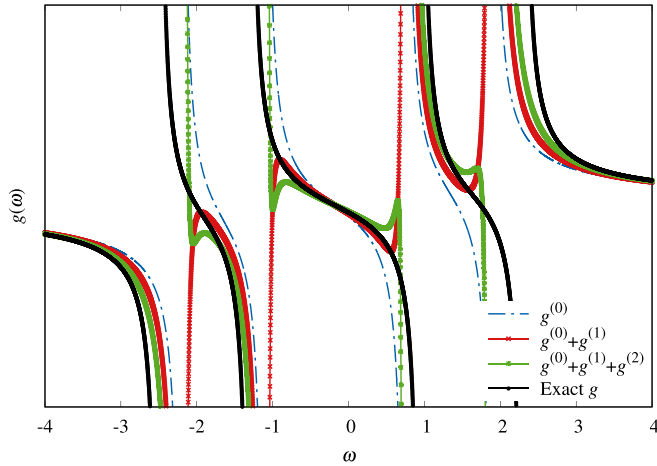


FIG. 23. Taylor expansions up to the second order in λ of the model Green's function $g(\omega)$ of Eq. (37).

where the near-vertical rapid oscillations of the nineteenth-order Taylor expansion occur.) In this overlap, both $g(\omega)$ and $g^{(0)}(\omega)$ are in the middle “shoulder” part of the monotonically decreasing functions of ω , displaying similar functional forms. Here, a Taylor expansion from $g^{(0)}(\omega)$ is convergent. Outside this overlap, either $g(\omega)$ or $g^{(0)}(\omega)$ is in the left “neck” part of the function, while the other is in the right “arm” part. They are both monotonically decreasing, but one of them has a singularity at one boundary of the domain, while the other has a singularity at the other boundary, and they have dissimilar functional forms in this sense. The $g(\omega)$ in this domain is nonanalytic and its Taylor expansion has zero radius of convergence.

In a molecular Green's function or self-energy, there are dense manifolds of singularities outside the central overlapping bracket, which encloses most principal roots and some low-energy satellite roots. Therefore, in practice, their perturbation expansions are safely convergent only in this central overlapping bracket as well as in the two terminal overlapping

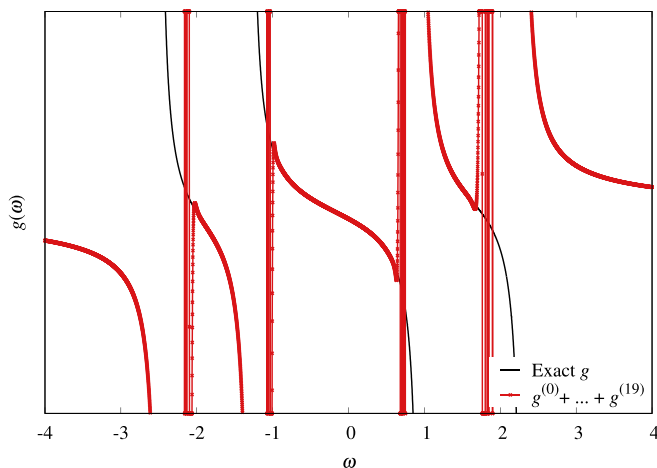


FIG. 24. Same as Fig. 23 but for the nineteenth-order Taylor expansion.

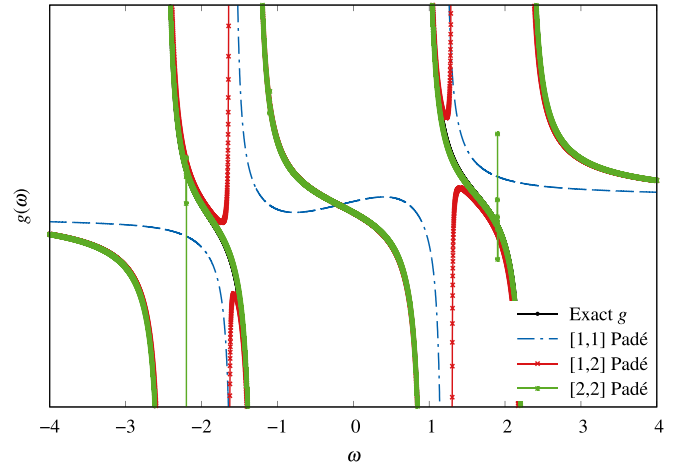


FIG. 25. Same as Fig. 23 but for Padé approximants.

brackets. The central overlapping bracket is an overlap of the bracket bounded by the highest 2p1h and lowest 2h1p HF orbital energy differences and the bracket demarcated typically by the least negative and least positive singularities of the exact self-energy [77].

B. Padé approximants

In Sec. V A, the nonconvergence is shown to be caused by the very definition of the exact Green's function, which is nonanalytic. Even when the positions of the poles, E_1, \dots, E_4 , are exactly expanded by second-order Taylor series, the Taylor expansion of $g(\omega)$ is not only inexact at the second order, but also nonconvergent at exactness at an infinite order. Therefore, the cause of nonconvergence is a mathematical one: The rational-function form of the exact Green's function is unamenable to a converging Taylor expansion, even though it contains full physical information about the poles.

For this mathematical problem, the purely mathematical resummation technique of Padé approximants [106] may prove useful. The $[M, N]$ Padé approximant of a function $f(\lambda)$ is a power-series expansion of a rational function and is defined by

$$[M, N]_f(\lambda) = \frac{a_0 + a_1\lambda + a_2\lambda^2 + \dots + a_M\lambda^M}{1 + b_1\lambda + b_2\lambda^2 + \dots + b_N\lambda^N}, \quad (43)$$

whose expansion coefficients, $\{a_0, \dots, a_M\}$ and $\{b_1, \dots, b_N\}$, are determined such that the above agrees with the $(M + N + 1)$ -order Taylor expansion of $f(\lambda)$. Padé approximants are known to generate a rapidly converging series out of even a divergent Taylor series, often working well “beyond their proven range of applicability” [106] at essentially no additional cost if the $(M + N + 1)$ -order Taylor expansion dominates the cost.

Figure 25 shows the $[1,1]$, $[1,2]$, and $[2,2]$ Padé approximants of $g(\omega)$. The $[2,2]$ Padé approximant (summing through the fifth-order Taylor series) is nearly exact and indistinguishable to the human eye from $g(\omega)$ except for a few spurious spikes. This is in contrast with Fig. 24 in which even the nineteenth-order Taylor expansion suffers from infinite errors in multiple ω domains. The remarkable performance of Padé

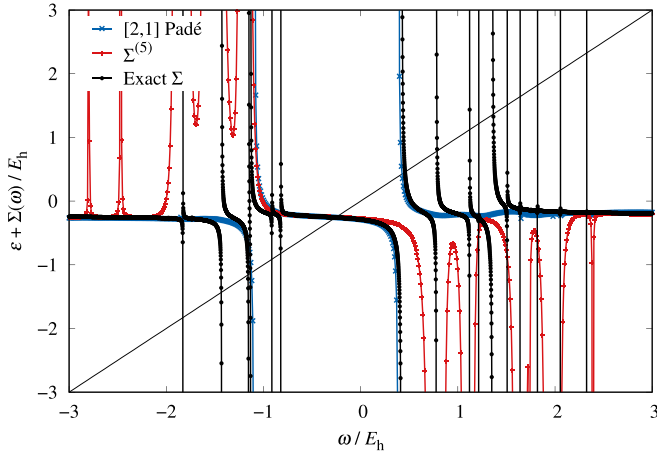


FIG. 26. The third diagonal element of $\epsilon + \Sigma(\omega)$ as a function of ω of the BH molecule obtained by a [2,1] Padé resummation of $\Sigma^{(n)}(\omega)$ over $1 \leq n \leq 5$. The corresponding elements of the exact $\epsilon + \Sigma(\omega)$ and $\epsilon + \Sigma^{(5)}(\omega)$ are also shown.

approximants is ascribed to separately expanding the numerator and denominator of the rational-function form of $g(\omega)$.

Padé approximants were introduced to many-body perturbation energies by Goscinski [99] and further developed by Brändas and Goscinski [100,102] and by Bartlett and Brändas [103,104]. They were applied to both convergent [105] and divergent [107] series of many-body perturbation energies, and in the latter case a rapid convergence was restored. In each case, the size-consistency is maintained [103,104]. Their applications to Green's functions were proposed by Goscinski and Lukman [34] and by Linderberg and Ratner [101], but no numerical results beyond a two-site model calculation were given.

Figures 26 and 27 show the [2,1] and [5,5] Padé approximants of the perturbative self-energies of the BH molecule. They were obtained by using Eqs. (12)–(17) of Laidig and Bartlett [105] of which their “ $E^{(n)}$ ” should be read $\Sigma^{(n)}$ in our case. The $[N, N - 1]$ and $[N, N]$ Padé approximants sum over perturbative self-energies through $2N + 1$ and $2N + 2$

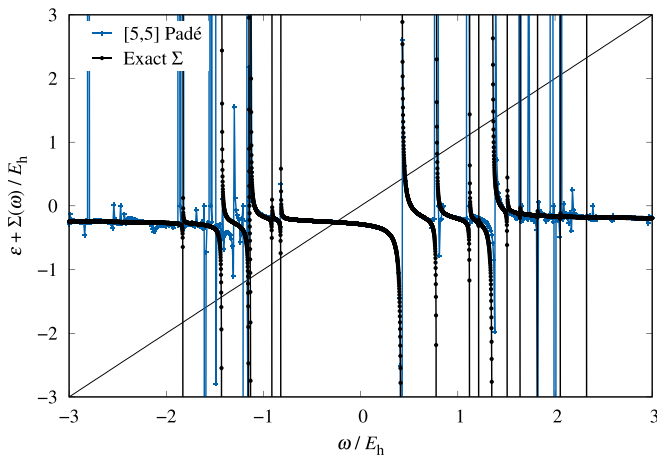


FIG. 27. The same as Fig. 26 but for a [5,5] Padé resummation of $\Sigma^{(n)}(\omega)$ over $1 \leq n \leq 12$.

orders, respectively, at no additional cost. The [2,1] Padé approximant (Fig. 26) not only largely eliminates the incorrect (concave and convex) shapes of $\Sigma^{(5)}$, but it also shifts the positions of the singularities in the right directions to be in better agreement with the exact self-energy. The [5,5] Padé approximant (Fig. 27) traces the exact self-energy even more closely, although spurious spikes can be seen.

In this context, it may be instructive to compare the ΔMPn method [36,63,64,162] with MBGF(n). ΔMPn computes the perturbative self-energy as the difference in the n th-order many-body perturbation energy between the frozen-orbital $N \pm 1$ and N electron systems. At the second and third orders, the ΔMPn self-energy agrees with the MBGF(n) self-energy in the diagonal, frequency-independent approximation [63,64]. At fourth and higher orders, these two self-energies deviate from each other, but they both converge at the same exact self-energy at an infinite order (unless they diverge). ΔMPn can reach the exact limit by including semireducible and linked-disconnected diagrams, which correct the errors from the diagonal and frequency-independent approximations, respectively. These strange diagrams (also implicit in EOM-CC) are, however, illegal in MBGF(n) [63,64].

Unlike MBGF(n), which is nonconvergent for most roots, ΔMPn is only occasionally divergent (see Fig. 6 of Ref. [64]) as it is based on the Hirschfelder-Certain degenerate perturbation theory (HCPT) [163] applied to individual states. This again underscores the fact that the positions of the poles can be directly and more reliably expanded in converging Taylor series; only when they are placed in the denominator of a rational function does a perturbation theory struggle to shift their positions in a systematic, converging manner. A Padé resummation of the perturbative self-energies works well because it expands the numerator and denominator of a Green's function separately and the power-series expansion of the denominator is analogous to ΔMPn .

VI. CONCLUSIONS

The Feynman-Dyson diagrammatic perturbation expansions of the self-energy and Green's function are reliably convergent and thus physically sound only in some small domains of ω , which includes the central overlapping bracket around $\omega = 0$ enclosing most principal roots and some low-energy satellite roots. Outside these domains, the perturbation expansions are nonconvergent. Our mathematical analysis suggests that the convergence is assured only in an overlap of the exact and mean-field ω brackets. An odd-order MBGF(n) method can have roots that are complex or whose residues are outside the valid range of zero to one, whereas a higher-even-order self-energy tends to consist of many vertical lines, predicting phantom poles with zero residues. A majority of satellite roots of MBGF(n) are distributed in nearly the same frequency domains spanned by mere HF orbital energy differences and include little to no electron-correlation effects, no matter how high the perturbation order is raised.

This is in contrast with the ΔMPn method [36,63,64,162], in which IPs and EAs are computed as the energy differences of the n th-order HCPT [163] applied to the frozen-orbital $N \pm 1$ and N electron systems. The HCPT energies for both principal and satellite ionized or electron-attached states [163]

are only occasionally divergent, and when they converge, they do so at the exact energies. Even though both ΔMPn and $MBGF(n)$ can be formulated in terms of the HCPT corrections to the identical zeroth-order wave functions and energies [64], the former is mostly sound for both principal and satellite states, while the latter is pathological for most satellite states.

Infinite partial resummations of diagrams can exacerbate the nonconvergence. The summation of the ring and ladder diagrams up to an infinite order by vertex renormalization (TDA) deteriorates the self-energy outside the central bracket. Worse still, the result of calculation changes dramatically and alternately with the number of cycles taken to solve the amplitude equations in an iterative algorithm, making the method ill-defined when all roots are sought. The summation of all tower diagrams by edge renormalization (self-consistent Green's function method) has a factorially increasing number of roots, whose energies encroach on the frequency domains where no exact roots can be found.

This is in sharp contrast with coupled-cluster theory. TDA is to IPs and EAs as LCCD is to ground-state energies since they both sum over the same infinite set of ring and ladder diagrams. Despite this similarity, TDA is pathological for low- and high-lying satellite roots, while coupled-cluster theory via the EOM-CC formalism offers the most accurate, robust, and converging approximations for all roots obtained directly as eigenvalues of a matrix [72,137–146]. The EOM formulation of MBGF [37,45–47,83], which can also be viewed as an infinite partial resummation of diagrams, is recast into a matrix diagonalization [48] and may thus be free from the convergence difficulties faced by $MBGF(n)$.

The nonconvergence of the Feynman-Dyson perturbation theory, which is the mathematical foundation of quantum field theory [20–24,27], makes it hard to use higher-than-second-order self-energy and Green's function in the *Ansätze* that are predicated on the knowledge of all poles and residues. Such *Ansätze* include the Galitskii-Migdal identity [108–111], self-consistent Green's function methods [8,18,88–98], and some models of the ADC [51,52] that evaluate the static part of its self-energy by summing over all poles of a Feynman-Dyson perturbative self-energy (see Deleuze *et al.* [112,113] for the ADC's lack of size- or charge-consistency, which may be related to the pathologies discussed here). Insofar as the self-consistent Green's function methods are derived from the diagrammatic Luttinger-Ward functional [8,15,16,19,88,89,114], on which the formalisms of DMFT [115] are also based, these methods may be adversely affected by the ill-defined nature of the perturbative Feynman propagators at many frequencies.

The nonconvergence is rooted in the rational-function form of the exact Green's function, featuring numerous singulari-

ties of the forms $(\omega - E_0 + E_I)^{-1}$ and $(\omega - E_A + E_0)^{-1}$ even before a perturbation approximation is introduced. This function is nonanalytic at many frequencies and thus cannot be expanded in converging power series, which is ultimately why perturbation theory tends to fail. This is reminiscent of the Bardeen-Cooper-Schrieffer theory of superconductivity [164], whose superconducting gap formula, $2\delta e^{-1/\rho V}$, is nonanalytic and cannot be expanded in a converging power series of the electron-phonon coupling V , even though it is small. To such a problem, “perturbation theory would not be easy to apply” (in page 224 of Ref. [25]). Likewise, the Kohn-Luttinger nonconvergence [2,8] of finite-temperature many-body perturbation theory [3–7] is ascribed to the non-analyticity of the rational-function forms of the exact grand potential and internal energy at $T = 0$ [9,10].

A Padé approximant is a power-series expansion of a rational function and is particularly promising for approximating a Feynman propagator, which is a nonanalytic rational function. Padé approximants were shown to accelerate the convergence of many-body perturbation energies, sometimes transforming a divergent series into a rapidly convergent one at no additional cost [99,100,102–105,107]. For $MBGF(n)$, Padé approximants exhibit remarkable performance, largely restoring the correct functional form of the self-energy and shift its poles in the right directions by the right amounts, which is consistent with our identified cause of the nonconvergence.

ACKNOWLEDGMENTS

We thank Prof. Piotr Piecuch (Michigan State University), Dr. Xiuyi Qin (CGG), Prof. Takashi Nakatsukasa (University of Tsukuba), and Prof. Vojtech Vlček (University of California, Santa Barbara) for insightful discussions and constructive suggestions. S.H. was supported by the U.S. Department of Energy (DoE), Office of Science, Office of Basic Energy Sciences under Grant No. DE-SC0006028 and also by the Center for Scalable Predictive methods for Excitations and Correlated phenomena (SPEC), which is funded by the U.S. DoE, Office of Science, Office of Basic Energy Sciences, Division of Chemical Sciences, Geosciences and Biosciences as part of the Computational Chemical Sciences (CCS) program at Pacific Northwest National Laboratory (PNNL) under FWP 70942. PNNL is a multiprogram national laboratory operated by Battelle Memorial Institute for the U.S. DoE. S.H. is a Guggenheim Fellow of the John Simon Guggenheim Memorial Foundation. I.G. was supported by Polish National Science Center under Grant No. 2020/39/O/ST4/00005. R.J.B. was supported by the Air Force Office of Scientific Research under AFOSR Award No. FA9550-19-1-0091.

-
- [1] F. J. Dyson, *Phys. Rev.* **85**, 631 (1952).
 [2] W. Kohn and J. M. Luttinger, *Phys. Rev.* **118**, 41 (1960).
 [3] P. K. Jha and S. Hirata, *Annu. Rep. Comput. Chem.* **15**, 3 (2019).
 [4] S. Hirata and P. K. Jha, *Annu. Rep. Comput. Chem.* **15**, 17 (2019).

- [5] S. Hirata and P. K. Jha, *J. Chem. Phys.* **153**, 014103 (2020).
 [6] P. K. Jha and S. Hirata, *Phys. Rev. E* **101**, 022106 (2020).
 [7] S. Hirata, *J. Chem. Phys.* **155**, 094106 (2021).
 [8] J. M. Luttinger and J. C. Ward, *Phys. Rev.* **118**, 1417 (1960).
 [9] S. Hirata, *Phys. Rev. A* **103**, 012223 (2021).
 [10] S. Hirata, *Chem. Phys. Lett.* **800**, 139668 (2022).

- [11] G. Lani, P. Romaniello, and L. Reining, *New J. Phys.* **14**, 013056 (2012).
- [12] T. Schäfer, G. Rohringer, O. Gunnarsson, S. Ciuchi, G. Sangiovanni, and A. Toschi, *Phys. Rev. Lett.* **110**, 246405 (2013).
- [13] V. Janiš and V. Pokorný, *Phys. Rev. B* **90**, 045143 (2014).
- [14] A. Stan, P. Romaniello, S. Rigamonti, L. Reining, and J. A. Berger, *New J. Phys.* **17**, 093045 (2015).
- [15] E. Kozik, M. Ferrero, and A. Georges, *Phys. Rev. Lett.* **114**, 156402 (2015).
- [16] R. Rossi and F. Werner, *J. Phys. A: Math. Theor.* **48**, 485202 (2015).
- [17] T. Schäfer, S. Ciuchi, M. Wallerberger, P. Thunström, O. Gunnarsson, G. Sangiovanni, G. Rohringer, and A. Toschi, *Phys. Rev. B* **94**, 235108 (2016).
- [18] W. Tarantino, P. Romaniello, J. A. Berger, and L. Reining, *Phys. Rev. B* **96**, 045124 (2017).
- [19] O. Gunnarsson, G. Rohringer, T. Schäfer, G. Sangiovanni, and A. Toschi, *Phys. Rev. Lett.* **119**, 056402 (2017).
- [20] R. P. Feynman, *Rev. Mod. Phys.* **20**, 367 (1948).
- [21] F. J. Dyson, *Phys. Rev.* **75**, 486 (1949).
- [22] F. J. Dyson, *Phys. Rev.* **75**, 1736 (1949).
- [23] J. Schwinger, *Proc. Natl. Acad. Sci. USA* **37**, 455 (1951).
- [24] J. J. Sakurai, *Advanced Quantum Mechanics* (Pearson, Reading, 1967).
- [25] N. H. March, W. H. Young, and S. Sampanthar, *The Many-Body Problem in Quantum Mechanics* (Cambridge University Press, Cambridge, 1967).
- [26] R. D. Mattuck, *A Guide to Feynman Diagrams in the Many-Body Problem* (Dover, New York, 1992).
- [27] F. Dyson, *Phys. World* **6**, 33 (1993).
- [28] A. L. Fetter and J. D. Walecka, *Quantum Theory of Many-Particle Systems* (Dover, New York, 2003).
- [29] L. P. Kadanoff and G. Baym, *Quantum Statistical Mechanics* (CRC Press, Boca Raton, 2018).
- [30] J. Lindenberg and Y. Öhrn, *Proc. Roy. Soc. A* **285**, 445 (1965).
- [31] L. Hedin, *Phys. Rev.* **139**, A796 (1965).
- [32] Y. Öhrn and J. Lindenberg, *Phys. Rev.* **139**, A1063 (1965).
- [33] J. Lindenberg and Y. Öhrn, *Chem. Phys. Lett.* **1**, 295 (1967).
- [34] O. Goscinski and B. Lukman, *Chem. Phys. Lett.* **7**, 573 (1970).
- [35] J. D. Doll and W. P. Reinhard, *J. Chem. Phys.* **57**, 1169 (1972).
- [36] B. T. Pickup and O. Goscinski, *Mol. Phys.* **26**, 1013 (1973).
- [37] J. Simons and W. D. Smith, *J. Chem. Phys.* **58**, 4899 (1973).
- [38] B. S. Yarlagadda, G. Csanak, H. S. Taylor, B. Schneider, and R. Yaris, *Phys. Rev. A* **7**, 146 (1973).
- [39] J. Lindenberg and Y. Öhrn, *Propagators in Quantum Chemistry* (Academic Press, London, 1973).
- [40] F. S. M. Tsui and K. F. Freed, *Chem. Phys.* **5**, 337 (1974).
- [41] J. Paldus and J. Čížek, *Adv. Quantum Chem.* **9**, 105 (1975).
- [42] L. S. Cederbaum, *J. Phys. B: At. Mol. Phys.* **8**, 290 (1975).
- [43] L. S. Cederbaum and W. Domcke, *Adv. Chem. Phys.* **36**, 205 (1977).
- [44] J. Simons, *Annu. Rev. Phys. Chem.* **28**, 15 (1977).
- [45] M. F. Herman, D. L. Yeager, and K. F. Freed, *Chem. Phys.* **29**, 77 (1978).
- [46] M. F. Herman, K. F. Freed, and D. L. Yeager, *J. Chem. Phys.* **72**, 602 (1980).
- [47] M. F. Herman, K. F. Freed, D. L. Yeager, and B. Liu, *J. Chem. Phys.* **72**, 611 (1980).
- [48] J. Baker and B. T. Pickup, *Chem. Phys. Lett.* **76**, 537 (1980).
- [49] Y. Öhrn and G. Born, *Adv. Quantum Chem.* **13**, 1 (1981).
- [50] P. Jørgensen and J. Simons, *Second Quantization-Based Methods in Quantum Chemistry* (Academic Press, New York, 1981).
- [51] J. Schirmer, *Phys. Rev. A* **26**, 2395 (1982).
- [52] J. Schirmer, L. S. Cederbaum, and O. Walter, *Phys. Rev. A* **28**, 1237 (1983).
- [53] W. von Niessen, J. Schirmer, and L. S. Cederbaum, *Comput. Phys. Rep.* **1**, 57 (1984).
- [54] M. D. Prasad, S. Pal, and D. Mukherjee, *Phys. Rev. A* **31**, 1287 (1985).
- [55] M. S. Hybertsen and S. G. Louie, *Phys. Rev. B* **34**, 5390 (1986).
- [56] J. Oddershede, Propagator methods, in *Advances in Chemical Physics: Ab Initio Methods in Quantum Chemistry Part 2*, edited by K. P. Lawley (Wiley, Hoboken, NJ, 1987), Vol. 69.
- [57] W. Kutzelnigg and D. Mukherjee, *J. Chem. Phys.* **107**, 432 (1997).
- [58] F. Aryasetiawan and O. Gunnarsson, *Rep. Prog. Phys.* **61**, 237 (1998).
- [59] J. V. Ortiz, *Adv. Quantum Chem.* **35**, 33 (1999).
- [60] G. Onida, L. Reining, and A. Rubio, *Rev. Mod. Phys.* **74**, 601 (2002).
- [61] J. V. Ortiz, *Wiley Interdiscip. Rev. Comput. Mol. Sci.* **3**, 123 (2013).
- [62] S. Y. Willow, K. S. Kim, and S. Hirata, *J. Chem. Phys.* **138**, 164111 (2013).
- [63] S. Hirata, M. R. Hermes, J. Simons, and J. V. Ortiz, *J. Chem. Theory Comput.* **11**, 1595 (2015).
- [64] S. Hirata, A. E. Doran, P. J. Knowles, and J. V. Ortiz, *J. Chem. Phys.* **147**, 044108 (2017).
- [65] C. M. Johnson, A. E. Doran, S. L. Ten-no, and S. Hirata, *J. Chem. Phys.* **149**, 174112 (2018).
- [66] A. E. Doran and S. Hirata, *J. Chem. Theory Comput.* **15**, 6097 (2019).
- [67] E. Opoku, F. Pawłowski, and J. V. Ortiz, *J. Chem. Phys.* **155**, 204107 (2021).
- [68] J. V. Ortiz, *Adv. Quantum Chem.* **85**, 109 (2022).
- [69] E. Opoku, F. Pawłowski, and J. V. Ortiz, *J. Chem. Theory Comput.* **18**, 4927 (2022).
- [70] E. Opoku, F. Pawłowski, and J. V. Ortiz, *J. Phys. Chem. A* **127**, 1085 (2023).
- [71] E. Opoku, F. Pawłowski, and J. V. Ortiz, *J. Chem. Phys.* **159**, 124109 (2023).
- [72] S. Hirata, M. Nooijen, and R. J. Bartlett, *Chem. Phys. Lett.* **328**, 459 (2000).
- [73] M. R. Hermes and S. Hirata, *J. Chem. Phys.* **139**, 034111 (2013).
- [74] X. Y. Qin and S. Hirata, *J. Phys. Chem. B* **124**, 10477 (2020).
- [75] S. Suhai, *Phys. Rev. B* **27**, 3506 (1983).
- [76] J.-Q. Sun and R. J. Bartlett, *Phys. Rev. Lett.* **77**, 3669 (1996).
- [77] J. Q. Sun and R. J. Bartlett, *J. Chem. Phys.* **104**, 8553 (1996).
- [78] S. Hirata and R. J. Bartlett, *J. Chem. Phys.* **112**, 7339 (2000).
- [79] S. Hirata and T. Shimazaki, *Phys. Rev. B* **80**, 085118 (2009).
- [80] S. Y. Willow, K. S. Kim, and S. Hirata, *Phys. Rev. B* **90**, 201110(R) (2014).
- [81] S. Hirata, Y. Shigeta, S. S. Xantheas, and R. J. Bartlett, *J. Phys. Chem. B* **127**, 3556 (2023).
- [82] B. D. Day, *Rev. Mod. Phys.* **39**, 719 (1967).
- [83] D. J. Rowe, *Rev. Mod. Phys.* **40**, 153 (1968).

- [84] M. Baldo, *Nuclear Methods and the Nuclear Equation of State* (World Scientific, Singapore, 1999).
- [85] G. D. Purvis and Y. Öhrn, *J. Chem. Phys.* **60**, 4063 (1974).
- [86] J. Schirmer and L. S. Cederbaum, *J. Phys. B: At. Mol. Phys.* **11**, 1889 (1978).
- [87] O. Walter and J. Schirmer, *J. Phys. B: At. Mol. Phys.* **14**, 3805 (1981).
- [88] G. Baym and L. P. Kadanoff, *Phys. Rev.* **124**, 287 (1961).
- [89] G. Baym, *Phys. Rev.* **127**, 1391 (1962).
- [90] D. Van Neck, M. Waroquier, and J. Ryckebusch, *Nucl. Phys. A* **530**, 347 (1991).
- [91] W. H. Dickhoff, The nucleon propagator in the nuclear medium, in *Nuclear Methods and the Nuclear Equation of State*, International Review of Nuclear Physics, edited by M. Baldo (World Scientific, Singapore, 1999), Vol. 8, Chap. 7, pp. 326–380.
- [92] W. H. Dickhoff and C. Barbieri, *Prog. Part. Nucl. Phys.* **52**, 377 (2004).
- [93] N. E. Dahlen and R. van Leeuwen, *J. Chem. Phys.* **122**, 164102 (2005).
- [94] C. Barbieri, *Phys. Lett. B* **643**, 268 (2006).
- [95] C. Barbieri and W. H. Dickhoff, *Int. J. Mod. Phys. A* **24**, 2060 (2009).
- [96] J. J. Phillips and D. Zgid, *J. Chem. Phys.* **140**, 241101 (2014).
- [97] D. Neuhauser, R. Baer, and D. Zgid, *J. Chem. Theory Comput.* **13**, 5396 (2017).
- [98] C. J. N. Coveney and D. P. Tew, *J. Chem. Theory Comput.* **19**, 3915 (2023).
- [99] O. Goscinski, *Int. J. Quantum Chem.* **1**, 769 (1967).
- [100] E. Brändas and O. Goscinski, *Phys. Rev. A* **1**, 552 (1970).
- [101] J. Linderberg and M. Ratner, *Chem. Phys. Lett.* **6**, 37 (1970).
- [102] O. Goscinski and E. Brändas, *Int. J. Quantum Chem.* **5**, 131 (1971).
- [103] R. J. Bartlett and E. J. Brändas, *J. Chem. Phys.* **56**, 5467 (1972).
- [104] R. J. Bartlett and E. J. Brändas, *J. Chem. Phys.* **59**, 2032 (1973).
- [105] W. D. Laidig, G. Fitzgerald, and R. J. Bartlett, *Chem. Phys. Lett.* **113**, 151 (1985).
- [106] C. M. Bender and S. A. Orszag, *Advanced Mathematical Methods for Scientists and Engineers I. Asymptotic Methods and Perturbation Theory* (Springer-Verlag, New York, 1999).
- [107] S. Hirata and R. J. Bartlett, *Chem. Phys. Lett.* **321**, 216 (2000).
- [108] V. M. Galitskii and A. B. Migdal, *Zh. Éxp. Teor. Fiz.* **34**, 139 (1958) [*Sov. Phys. JETP* **7**, 96 (1958)].
- [109] D. S. Koltun, *Phys. Rev. Lett.* **28**, 182 (1972).
- [110] L. J. Holleboom and J. G. Snijders, *J. Chem. Phys.* **93**, 5826 (1990).
- [111] J. V. Ortiz, *Int. J. Quantum Chem.* **56**, 331 (1995).
- [112] M. Deleuze, M. K. Scheller, and L. S. Cederbaum, *J. Chem. Phys.* **103**, 3578 (1995).
- [113] M. S. Deleuze, *Int. J. Quantum Chem.* **93**, 191 (2003).
- [114] L. Lin and M. Lindsey, *Proc. Natl. Acad. Sci. USA* **115**, 2282 (2018).
- [115] A. Georges, G. Kotliar, W. Krauth, and M. J. Rozenberg, *Rev. Mod. Phys.* **68**, 13 (1996).
- [116] I. Shavitt and R. J. Bartlett, *Many-Body Methods in Chemistry and Physics* (Cambridge University Press, Cambridge, 2009).
- [117] J. V. Ortiz, *J. Chem. Phys.* **153**, 070902 (2020).
- [118] R. J. Bartlett, I. Grabowski, S. Hirata, and S. Ivanov, *J. Chem. Phys.* **122**, 034104 (2005).
- [119] R. J. Bartlett, *Chem. Phys. Lett.* **484**, 1 (2009).
- [120] R. Manne and T. Åberg, *Chem. Phys. Lett.* **7**, 282 (1970).
- [121] A. Szabo and N. S. Ostlund, *Modern Quantum Chemistry* (MacMillan, New York, 1982).
- [122] O. Walter, L. S. Cederbaum, and J. Schirmer, *J. Math. Phys.* **25**, 729 (1984).
- [123] P. Piecuch, S. Zarrabian, J. Paldus, and J. Čížek, *Phys. Rev. A* **42**, 5155 (1990).
- [124] S. Hirata, *Theor. Chem. Acta* **129**, 727 (2011).
- [125] V. Vlček, *J. Chem. Theory Comput.* **15**, 6254 (2019).
- [126] C. Mejuto-Zaera and V. Vlček, *Phys. Rev. B* **106**, 165129 (2022).
- [127] R. J. Bartlett and M. Musiał, *Rev. Mod. Phys.* **79**, 291 (2007).
- [128] J. Čížek, *J. Chem. Phys.* **45**, 4256 (1966).
- [129] A. C. Hurley, *Electron Correlation in Small Molecules* (Academic, New York, 1976).
- [130] R. J. Bartlett and I. Shavitt, *Chem. Phys. Lett.* **50**, 190 (1977).
- [131] R. J. Bartlett and I. Shavitt, *Chem. Phys. Lett.* **57**, 157 (1978).
- [132] R. J. Bartlett, I. Shavitt, and G. D. Purvis, *J. Chem. Phys.* **71**, 281 (1979).
- [133] R. Ahlrichs, *Comput. Phys. Commun.* **17**, 31 (1979).
- [134] S. Koch and W. Kutzelnigg, *Theor. Chim. Acta* **59**, 387 (1981).
- [135] A. G. Taube and R. J. Bartlett, *J. Chem. Phys.* **130**, 144112 (2009).
- [136] E. Bergli and M. Hjorth-Jensen, *Ann. Phys. (NY)* **326**, 1125 (2011).
- [137] J. F. Stanton and R. J. Bartlett, *J. Chem. Phys.* **98**, 7029 (1993).
- [138] J. F. Stanton and J. Gauss, *J. Chem. Phys.* **101**, 8938 (1994).
- [139] M. Nooijen and R. J. Bartlett, *J. Chem. Phys.* **102**, 6735 (1995).
- [140] R. J. Bartlett, J. E. DelBene, S. A. Perera, and R. P. Mattie, *J. Mol. Struct.: THEOCHEM* **400**, 157 (1997).
- [141] J. F. Stanton and J. Gauss, *J. Chem. Phys.* **111**, 8785 (1999).
- [142] J. R. Gour, P. Piecuch, and M. Włoch, *J. Chem. Phys.* **123**, 134113 (2005).
- [143] M. Kamiya and S. Hirata, *J. Chem. Phys.* **125**, 074111 (2006).
- [144] J. R. Gour and P. Piecuch, *J. Chem. Phys.* **125**, 234107 (2006).
- [145] M. Kamiya and S. Hirata, *J. Chem. Phys.* **126**, 134112 (2007).
- [146] R. J. Bartlett, *Wiley Interdiscip. Rev. Comput. Mol. Sci.* **2**, 126 (2012).
- [147] K. Schönhammer and O. Gunnarsson, *Phys. Rev. B* **18**, 6606 (1978).
- [148] M. Nooijen and J. G. Snijders, *Int. J. Quantum Chem.* **44**, 55 (1992).
- [149] M. Nooijen and J. G. Snijders, *Int. J. Quantum Chem.* **48**, 15 (1993).
- [150] L. Meissner and R. J. Bartlett, *Int. J. Quantum Chem.* **48**, 67 (1993).
- [151] K. Kowalski, K. Bhaskaran-Nair, and W. A. Shelton, *J. Chem. Phys.* **141**, 094102 (2014).
- [152] B. Peng and K. Kowalski, *Phys. Rev. A* **94**, 062512 (2016).
- [153] K. Bhaskaran-Nair, K. Kowalski, and W. A. Shelton, *J. Chem. Phys.* **144**, 144101 (2016).
- [154] B. Peng, R. Van Beeumen, D. B. Williams-Young, K. Kowalski, and C. Yang, *J. Chem. Theory Comput.* **15**, 3185 (2019).

- [155] B. Peng, K. Kowalski, A. Panyala, and S. Krishnamoorthy, *J. Chem. Phys.* **152**, 011101 (2020).
- [156] J. J. Rehr, F. D. Vila, J. J. Kas, N. Y. Hirshberg, K. Kowalski, and B. Peng, *J. Chem. Phys.* **152**, 174113 (2020).
- [157] H. Pathak, A. Panyala, B. Peng, N. P. Bauman, E. Mutlu, J. J. Rehr, F. D. Vila, and K. Kowalski, *J. Chem. Theory Comput.* **19**, 2248 (2023).
- [158] M. L. Vidal, X. T. Feng, E. Epifanovsky, A. I. Krylov, and S. Coriani, *J. Chem. Theory Comput.* **15**, 3117 (2019).
- [159] P. R. Surján and A. Szabados, *Int. J. Quantum Chem.* **69**, 713 (1998).
- [160] A. A. Kananenka, A. R. Welden, T. N. Lan, E. Gull, and D. Zgid, *J. Chem. Theory Comput.* **12**, 2250 (2016).
- [161] T. N. Lan, A. A. Kananenka, and D. Zgid, *J. Chem. Theory Comput.* **12**, 4856 (2016).
- [162] S. Šmiga and I. Grabowski, *J. Chem. Theory Comput.* **14**, 4780 (2018).
- [163] J. O. Hirschfelder and P. R. Certain, *J. Chem. Phys.* **60**, 1118 (1974).
- [164] J. Bardeen, L. N. Cooper, and J. R. Schrieffer, *Phys. Rev.* **108**, 1175 (1957).