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Abstract
A Görling–Levy (GL)-based perturbation theory along the range-separated adiabatic connection is assessed for the calculation of electronic excitation energies. In comparison with the Rayleigh–Schrödinger (RS)-based perturbation theory introduced in a previous work [E. Rebolini, J. Toulouse, A. M. Teale, T. Helgaker, A. Savin, Mol. Phys. **113**, 1740 (2015)], this GL-based perturbation theory keeps the ground-state density constant at each order and thus gives the correct ionization energy at each order. Excitation energies up to first order in the perturbation have been calculated numerically for the helium and beryllium atoms and the hydrogen molecule without introducing any density-functional approximations. In comparison with the RS-based perturbation theory, the present GL-based perturbation theory gives much more accurate excitation energies for Rydberg states but similar excitation energies for valence states.

KEYWORDS
density-functional theory; range separation; adiabatic connection; perturbation theory; excitation energies

1. Introduction

Within the framework of density-functional theory (DFT), the calculation of molecular excitation energies is nowadays mostly performed using linear-response time-dependent density-functional theory (TDDFT) (see, e.g., Refs. [1, 2]) within the adiabatic local or semi-local approximations. Despite its many successes, linear-response

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TDDFT within these approximations suffers from serious limitations, especially for describing systems with static (or strong) correlation [3], double or multiple excitations [4], and Rydberg and charge-transfer excitations [5, 6]. These deficiencies have been attributed to the locality of the approximated exchange–correlation potential and kernel either in space (local and semi-local approximations) or in time (adiabatic approximation). While the former is directly linked to functional development in time-independent DFT, the latter is a problem specific to the time-dependent formulation. However, time dependence is in principle not required to describe excited states since by the Hohenberg–Kohn theorem [7] the time-independent ground-state density contains all the information about the system including information about its excited states.

Over the years, several time-independent DFT approaches for calculating excitation energies have emerged and are still being developed: ensemble DFT [8–17], state-specific self-consistent DFT and related methods [18–39], hybrid DFT/configuration interaction (CI) methods [40–44] and perturbation theory starting from the non-interacting Kohn–Sham (KS) Hamiltonian [45–48]. In a previous work [49], we have explored further this density-functional perturbation-theory approach with one key modification: As a zeroth-order Hamiltonian, instead of using the non-interacting KS Hamiltonian, we use a *partially interacting Hamiltonian* incorporating the *long-range* part only of the Coulomb electron–electron interaction, corresponding to an intermediate point along a range-separated adiabatic connection [50–56]. The partially interacting zeroth-order Hamiltonian is of course closer to the exact Hamiltonian than is the non-interacting KS Hamiltonian, thereby putting less demand on the perturbation theory.

In this previous work [49], a Rayleigh–Schrödinger (RS)-based perturbation theory was tested on a few atoms and molecules and it was found that the first-order excitation energies are not overall improved in comparison with the zeroth-order excitation energies. This finding was rationalized by the fact that this perturbation theory does not keep the ground-state density constant at each order. In the present work, we explore an alternative approach, based on Görling–Levy (GL) perturbation theory along a range-separated adiabatic connection, which keeps the ground-state density constant at each order and therefore should give more accurate excitation energies.

The paper is organized as follows. The main equations of our GL-based range-separated perturbation theory are given in Section 2. The computational details for the calculations carried out, involving no other approximations than the use of a finite basis, can be found in Section 3. The results obtained for the He and Be atoms and for the H_2 molecule are discussed in Section 4. Finally, Section 5 contains our conclusions.

2. Theoretical background

This section consists of two parts. We first review range-separated DFT for ground states in Subsection 2.1 and then GL-based perturbation theory for excited states in Subsection 2.2.

2.1. *Range-separated ground-state density-functional theory*

In range-separated DFT (see, e.g., Ref. [54]), the electron–electron interaction is partitioned into long-range and short-range contributions by means of the error function and of a range-separation parameter μ which controls the range of the separation. The

long-range (lr) interaction operator is defined as

$$\hat{W}_{\rm ee}^{\rm lr,\mu} = \frac{1}{2} \iint w_{\rm ee}^{\rm lr,\mu}(r_{12}) \hat{n}_2(\mathbf{r}_1, \mathbf{r}_2) \mathrm{d}\mathbf{r}_1 \mathrm{d}\mathbf{r}_2, \tag{1}$$

and is written in terms of the error-function interaction $w_{\rm ee}^{\rm lr,\mu}(r) = \mathrm{erf}(\mu r)/r$ and the pair-density operator $\hat{n}_2(\mathbf{r}_1, \mathbf{r}_2)$, where $\mathbf{r}$ refers to the electron coordinates. The exact ground-state energy of an N-electron system is then expressed as

$$E_0 = \min_{\Psi} \Big\{ \langle \Psi | \hat{T} + \hat{V}_{\rm ne} + \hat{W}_{\rm ee}^{\rm lr,\mu} | \Psi \rangle + \bar{E}_{\rm Hxc}^{\rm sr,\mu}[n_\Psi] \Big\}, \tag{2}$$

where the minimization is performed over normalized multi-determinantal wave functions Ψ. In Eq. (2), we have introduced the kinetic-energy operator $\hat{T}$, the nuclear–electron attraction operator $\hat{V}_{\rm ne} = \int v_{\rm ne}(\mathbf{r}) \hat{n}(\mathbf{r}) \mathrm{d}\mathbf{r}$ written in terms of the density operator $\hat{n}(\mathbf{r})$, and the complement short-range (sr) Hartree–exchange–correlation density functional $\bar{E}_{\rm Hxc}^{\rm sr,\mu}[n_\Psi]$ evaluated at the density of Ψ, $n_\Psi(\mathbf{r}) = \langle \Psi | \hat{n}(\mathbf{r}) | \Psi \rangle$. The minimizing wave function Ψ_0^μ in Eq. (2) is the ground-state wave function of the following eigenvalue problem

$$\hat{H}^{\rm lr,\mu} | \Psi_0^\mu \rangle = \mathcal{E}_0^\mu | \Psi_0^\mu \rangle, \tag{3}$$

associated with the energy $\mathcal{E}_0^\mu$. In Eq. (3), we have introduced the partially interacting Hamiltonian

$$\hat{H}^{\rm lr,\mu} = \hat{T} + \hat{V}_{\rm ne} + \hat{W}_{\rm ee}^{\rm lr,\mu} + \hat{\bar{V}}_{\rm Hxc}^{\rm sr,\mu}, \tag{4}$$

which contains the operator

$$\hat{\bar{V}}_{\rm Hxc}^{\rm sr,\mu} = \int \bar{v}_{\rm Hxc}^{\rm sr,\mu}(\mathbf{r}) \hat{n}(\mathbf{r}) \mathrm{d}\mathbf{r}, \tag{5}$$

with the short-range Hartree–exchange–correlation potential $\bar{v}_{\rm Hxc}^{\rm sr,\mu}(\mathbf{r}) = \delta \bar{E}_{\rm Hxc}^{\rm sr,\mu}[n_0]/\delta n(\mathbf{r})$ evaluated at the exact ground-state density $n_0(\mathbf{r})$. This is the potential that keeps the ground-state density constant for all μ, i.e. $\langle \Psi_0^\mu | \hat{n}(\mathbf{r}) | \Psi_0^\mu \rangle = n_0(\mathbf{r})$.

At $\mu = 0$, the Hamiltonian $\hat{H}^{\rm lr,\mu}$ in Eq. (4) reduces to the standard non-interacting KS Hamiltonian, whereas for $\mu \to \infty$ it reduces to the physical Hamiltonian. The Hamiltonian $\hat{H}^{\rm lr,\mu}$ therefore defines a range-separated adiabatic connection, linking the KS and the physical systems when varying μ, keeping the ground-state density constant.

2.2. *Excited states from GL-based perturbation theory*

For a given value of μ, the excited-state wave functions and energies of the long-range interacting Hamiltonian

$$\hat{H}^{\rm lr,\mu} | \Psi_k^\mu \rangle = \mathcal{E}_k^\mu | \Psi_k^\mu \rangle, \tag{6}$$

can be used as zeroth-order approximations to the physical excited-state wave functions and energies. It can then be improved upon by setting up perturbation theories

with $\hat{H}^{\mathrm{lr},\mu}$ as the zeroth-order Hamiltonian. As shown in Ref. [49], an RS-based perturbation theory in which the ground-state density is not kept constant gives first-order excitation energies that overall do not improve upon the zeroth-order excitation energies. Here we instead use a GL-based perturbation theory, with the ground-state density kept constant.

To formulate such a GL-based perturbation theory, we define the following Hamiltonian depending on a coupling constant λ

$$\hat{H}^{\mu,\lambda} = \hat{T} + \hat{V}_{\mathrm{ne}} + \hat{W}_{\mathrm{ee}}^{\mathrm{lr},\mu} + \lambda \hat{W}_{\mathrm{ee}}^{\mathrm{sr},\mu} + \hat{\bar{V}}_{\mathrm{Hxc}}^{\mathrm{sr},\mu,\lambda}, \tag{7}$$

which contains the operator

$$\hat{\bar{V}}_{\mathrm{Hxc}}^{\mathrm{sr},\mu,\lambda} = \int \bar{v}_{\mathrm{Hxc}}^{\mathrm{sr},\mu,\lambda}(\mathbf{r}) \hat{n}(\mathbf{r}) \, \mathrm{d}\mathbf{r}, \tag{8}$$

where $\bar{v}_{\mathrm{Hxc}}^{\mathrm{sr},\mu,\lambda}(\mathbf{r})$ is the λ-dependent short-range Hartree–exchange–correlation potential that keeps the ground-state density constant for all μ and all λ – that is, $\langle \Psi_0^{\mu,\lambda} | \hat{n}(\mathbf{r}) | \Psi_0^{\mu,\lambda} \rangle = n_0(\mathbf{r})$ where $\Psi_0^{\mu,\lambda}$ is the ground-state wave function of the Hamiltonian $\hat{H}^{\mu,\lambda}$ in Eq. (7). The Hamiltonian $\hat{H}^{\mu,\lambda}$ thus sets up a double adiabatic connection with a constant ground-state density. To clearly separate the linear and non-linear dependence in λ, we then rewrite Eq. (7) as

$$\hat{H}^{\mu,\lambda} = \hat{T} + \hat{V}_{\mathrm{ne}} + \hat{W}_{\mathrm{ee}}^{\mathrm{lr},\mu} + \hat{\bar{V}}_{\mathrm{Hxc}}^{\mathrm{sr},\mu} + \lambda \hat{W}_{\mathrm{ee}}^{\mathrm{sr},\mu} + \hat{\bar{V}}_{\mathrm{Hxc}}^{\mathrm{sr},\mu,\lambda} - \hat{\bar{V}}_{\mathrm{Hxc}}^{\mathrm{sr},\mu}, \tag{9}$$

and we define the potential operator

$$\hat{V}_{\mathrm{Hxc}}^{\mathrm{sr},\mu,\lambda} = \hat{\bar{V}}_{\mathrm{Hxc}}^{\mathrm{sr},\mu} - \hat{\bar{V}}_{\mathrm{Hxc}}^{\mathrm{sr},\mu,\lambda}, \tag{10}$$

which can be decomposed into a linear contribution with respect to λ and a term containing all higher-order terms (see Ref. [49])

$$\hat{V}_{\mathrm{Hxc}}^{\mathrm{sr},\mu,\lambda} = \lambda \hat{V}_{\mathrm{Hx,md}}^{\mathrm{sr},\mu} + \hat{V}_{\mathrm{c,md}}^{\mathrm{sr},\mu,\lambda}. \tag{11}$$

The potential $\hat{V}_{\mathrm{Hx,md}}^{\mathrm{sr},\mu}$ is the short-range "multi-determinantal (md) Hartree–exchange" potential operator, while $\hat{V}_{\mathrm{c,md}}^{\mathrm{sr},\mu,\lambda}$ is the short-range "multi-determinantal correlation" potential operator [57]. For non-degenerate ground-state wave functions Ψ_0^{μ}, the expansion of $\hat{V}_{\mathrm{c,md}}^{\mathrm{sr},\mu,\lambda}$ in λ around $\lambda = 0$ begins at second order:

$$\hat{V}_{\mathrm{c,md}}^{\mathrm{sr},\mu,\lambda} = \lambda^2 \, \hat{V}_{\mathrm{c,md}}^{\mathrm{sr},\mu,(2)} + \lambda^3 \, \hat{V}_{\mathrm{c,md}}^{\mathrm{sr},\mu,(3)} + \cdots . \tag{12}$$

The partially interacting Hamiltonian can then be rewritten as

$$\begin{aligned} \hat{H}^{\mu,\lambda} &= \hat{H}^{\mathrm{lr},\mu} + \lambda \hat{W}^{\mathrm{sr},\mu} - \hat{V}_{\mathrm{c,md}}^{\mathrm{sr},\mu,\lambda}, \\ &= \hat{H}^{\mathrm{lr},\mu} + \lambda \hat{W}^{\mathrm{sr},\mu} - \lambda^2 \, \hat{V}_{\mathrm{c,md}}^{\mathrm{sr},\mu,(2)} - \lambda^3 \, \hat{V}_{\mathrm{c,md}}^{\mathrm{sr},\mu,(3)} + \cdots , \end{aligned} \tag{13}$$

where the unperturbed Hamiltonian is the partially interacting Hamiltonian defined in Eq. (4) and $\hat{W}^{\mathrm{sr},\mu} = \hat{W}_{\mathrm{ee}}^{\mathrm{sr},\mu} - \hat{V}_{\mathrm{Hx,md}}^{\mathrm{sr},\mu}$ is the perturbation operator. More details on

this GL-based perturbation theory can be found in the Appendix of Ref. [49].

The excited-state wave functions and energies of the perturbed Hamiltonian are thus expanded with respect to λ as

$$|\Psi_k^{\mu,\lambda}\rangle = |\Psi_k^{\mu}\rangle + \lambda|\Psi_k^{\mu,(1)}\rangle + \lambda^2|\Psi_k^{\mu,(2)}\rangle + \lambda^3|\Psi_k^{\mu,(3)}\rangle + \cdots \tag{14}$$

$$E_k^{\mu,\lambda} = \mathcal{E}_k^{\mu} + \lambda E_k^{\mu,(1)} + \lambda^2 E_k^{\mu,(2)} + \lambda^3 E_k^{\mu,(3)} + \cdots \tag{15}$$

where $\Psi_k^{\mu,(n)}$ and $E_k^{\mu,(n)}$ are the n-th order corrections to the wave function and energy of the k-th state. In particular, the first-order energy correction is

$$E_k^{\mu,(1)} = \langle \Psi_k^{\mu}|\hat{W}^{\mathrm{sr},\mu}|\Psi_k^{\mu}\rangle, \tag{16}$$

and the corresponding zeroth+first order energy is

$$E_k^{\mu,(0+1)} = \mathcal{E}_k^{\mu} + E_k^{\mu,(1)} = \langle \Psi_k^{\mu}|\hat{H} + \hat{\bar{V}}_{\mathrm{c,md}}^{\mathrm{sr},\mu}|\Psi_k^{\mu}\rangle. \tag{17}$$

Equation (17) contains two contributions; the zeroth+first order energy in standard RS perturbation theory, $E_{k,\mathrm{RS}}^{\mu,(0+1)} = \langle \Psi_k^{\mu}|\hat{H}|\Psi_k^{\mu}\rangle$, and an additional term, $\langle \Psi_k^{\mu}|\hat{\bar{V}}_{\mathrm{c,md}}^{\mathrm{sr},\mu}|\Psi_k^{\mu}\rangle$, which is only present in GL-based perturbation theory.

A local-density approximation has been developed for the short-range multideterminant correlation energy functional (and the associated potential in $\hat{\bar{V}}_{\mathrm{c,md}}^{\mathrm{sr},\mu}$) [57, 58]. In the present study, we will test the GL-based perturbation theory without introducing any density-functional approximations, providing benchmark data for future approximations.

3. Computational details

Calculations were performed for the He and Be atoms and the H_2 molecule with a development version of the DALTON program [59, 60]; see Refs. [55, 61, 62]. Following the same procedure as in Ref. [56], a full CI (FCI) calculation was first carried out to obtain the exact ground-state density in the chosen basis set. Next, for several values of μ and λ, a Lieb optimization [61, 63, 64] was carried out to obtain the short-range potential $\bar{v}^{\mathrm{sr},\mu,\lambda}(\mathbf{r}) = v_{\mathrm{ne}}(\mathbf{r}) + \bar{v}_{\mathrm{Hxc}}^{\mathrm{sr},\mu,\lambda}(\mathbf{r})$ needed in Eq. (7) to reproduce this FCI ground-state density with the partial electron–electron interaction $w_{\mathrm{ee}}^{\mathrm{lr},\mu}(r_{12}) + \lambda w_{\mathrm{ee}}^{\mathrm{sr},\mu}(r_{12})$. Then, an FCI calculation was performed with the partially-interacting Hamiltonian constructed from $w_{\mathrm{ee}}^{\mathrm{lr},\mu}(r_{12}) + \lambda w_{\mathrm{ee}}^{\mathrm{sr},\mu}(r_{12})$ and $\bar{v}^{\mathrm{sr},\mu,\lambda}(\mathbf{r})$ to obtain, for a few states, the energies $E_k^{\mu,\lambda}$ and wave functions $\Psi_k^{\mu,\lambda}$, for each values of μ and λ.

For each system, each excited state, and each value of μ, a third-degree polynomial fit in λ was performed on the excitation energy $\Delta E_k^{\mu,\lambda} = E_k^{\mu,\lambda} - E_0^{\mu,\lambda}$ for values of λ ranging from 0 to 0.3 and the first-order correction in the GL-based perturbation theory was obtained as the first-order derivative with respect to λ. The zeroth-order excitation energies and the first-order excitation energies in the RS-based perturbation theory were already available from Refs. [56] and [49], respectively.

The basis sets used were uncontracted t-aug-cc-pV5Z for He, uncontracted d-aug-cc-pVDZ for Be, and uncontracted d-aug-cc-pVTZ for H_2.

4. Results and discussion

4.1. *First Rydberg excitation energies of the helium atom*

The first singlet and triplet excitation energies of the helium atom correct to zeroth and first orders along the range-separated adiabatic connection in the GL- and RS-based perturbation theories are shown in Figure 1.

In the KS limit, at $\mu = 0$, the zeroth-order singlet and triplet excitation energies are degenerate. When increasing μ, this degeneracy is lifted and the zeroth-order singlet and triplet excitation energies eventually converge to their physical excitation energies for $\mu \to \infty$. With the introduction of the perturbation, the singlet/triplet degeneracy is lifted already at $\mu = 0$. As found in Ref. [49], the first-order RS-based perturbative correction globally deteriorates the zeroth-order excitation energies, leading to large errors that do not decrease monotonically with μ.

Interestingly, the zeroth+first order excitation energies obtained from the GL-based perturbation theory have much smaller errors than those obtained by the RS-based perturbation theory. In particular, the first-order GL correction systematically improves the zeroth-order excitation energies for sufficiently large μ, accelerating the convergence with respect to the range separation parameter to the physical excitation energies: a 1 millihartree accuracy is reached for μ larger than 1.2 bohr^{-1}, while values of 3 to 4 bohr^{-1} are necessary for the zeroth-order curves. However, near $\mu = 0$, the first-order GL correction does not always improve the zeroth-order excitation energy (see the $1^1\mathrm{S} \to 1^1\mathrm{P}$ transition).

These comparisons show that the GL-based perturbation theory (which keeps the ground-state density constant, ensuring a correct ionization energy) is a much better strategy for calculating Rydberg excitation energies than the RS-based perturbation theory. This was expected since these Rydberg excitation energies are close to the ionization threshold.

4.2. *Valence excitation energies of the beryllium atom*

The valence excitation energies of the beryllium atom correct to zeroth and first orders in the GL-based and RS-based perturbation theories are plotted in Figure 2. For these excitations, the two approaches give very similar first-order corrections. This behaviour can be rationalized from the fact that these valence excited states are far from the ionization threshold so that imposing the correct ionization energy (by keeping the ground-state density constant) has much less impact than for the Rydberg excitation energies.

We note that the first-order correction systematically improves the singlet excitation energy but not the triplet excitation energy, which is overestimated at zeroth order and underestimated by about the same amount at first order.

4.3. *First valence excitation energies of the hydrogen molecule*

In Figure 3, we have plotted the first valence excitation energies of hydrogen molecule at its equilibrium internuclear distance R_{eq} and at $3R_{\mathrm{eq}}$. As for the valence excitation energies in beryllium, the first-order RS- and GL-based corrections are overall quite similar, but with a discernible faster μ-convergence of the GL-based zeroth+first order excitation energies to the physical excitation energies at the equilibrium distance.

At the equilibrium distance, the GL-based first-order perturbation theory overshoots

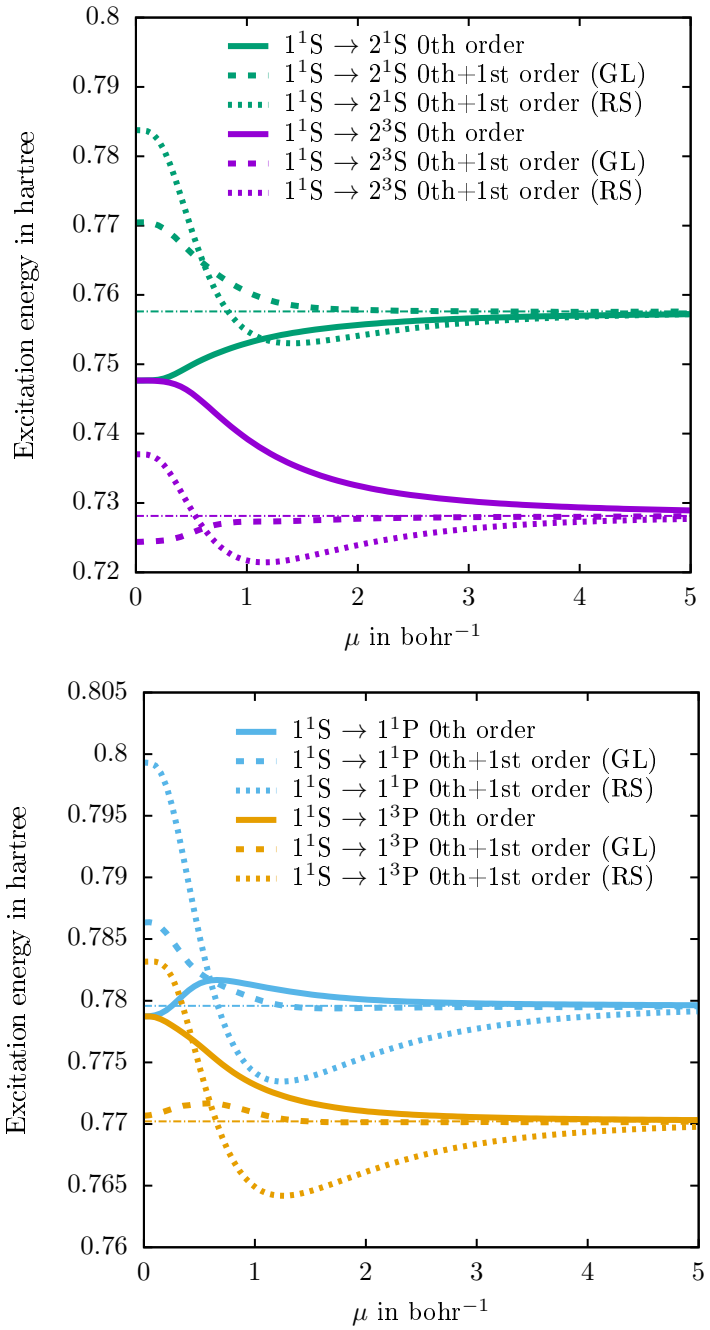

Figure 1. Zeroth-order excitation energies $\Delta\mathcal{E}_k^\mu = \mathcal{E}_k^\mu - \mathcal{E}_0^\mu$ (plain line) and zeroth+first order excitation energies $\Delta E_k^{\mu,(0+1)} = E_k^{\mu,(0+1)} - E_0^{\mu,(0+1)}$ (dashed line) in the GL-based perturbation theory for the helium atom as a function of μ. For comparison, the zeroth+first order excitation energies $\Delta E_{k,\mathrm{RS}}^{\mu,(0+1)} = E_{k,\mathrm{RS}}^{\mu,(0+1)} - E_{0,\mathrm{RS}}^{\mu,(0+1)}$ (dotted line) in the RS-based perturbation theory of Ref. [49] are also shown. The excitation energies of the physical system $\Delta E_k = E_k - E_0$ are plotted as thin horizontal dot-dashed lines.

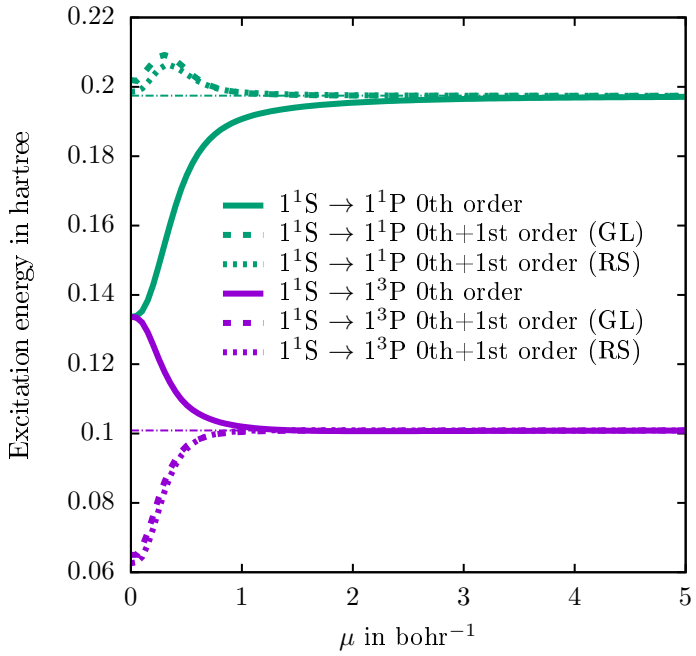

Figure 2. Zeroth-order excitation energies $\Delta\mathcal{E}_k^\mu = \mathcal{E}_k^\mu - \mathcal{E}_0^\mu$ (plain line) and zeroth+first order excitation energies $\Delta E_k^{\mu,(0+1)} = E_k^{\mu,(0+1)} - E_0^{\mu,(0+1)}$ (dashed line) in the GL-based perturbation theory for the beryllium atom as a function of μ. For comparison, the zeroth+first order excitation energies $\Delta E_{k,\mathrm{RS}}^{\mu,(0+1)} = E_{k,\mathrm{RS}}^{\mu,(0+1)} - E_{0,\mathrm{RS}}^{\mu,(0+1)}$ (dotted line) in the RS-based perturbation theory of Ref. [49] are also shown. The excitation energies of the physical system $\Delta E_k = E_k - E_0$ are plotted as thin horizontal dot-dashed lines.

the correction to both the singlet and triplet zeroth-order excitation energies for small values of μ, but nevertheless improves upon the zero-order correction for all μ values.

When the bond is stretched, the first-order correction no longer systematically improves on the zeroth-order excitation energies for small μ. In particular, the excitation energy of the first transition $1^1\Sigma_\mathrm{g}^+ \to 1^3\Sigma_\mathrm{u}^+$ becomes negative for small μ and the error with respect to the physical excitation energy is higher than in the zeroth-order case. Moreover, the ordering of the two excitation energies for the transitions to the two singlet states is incorrect at small μ. For values of μ greater than about 1 bohr^{-1}, the first-order correction does provide a systematic improvement over the zeroth-order excitation energies, particularly for the transition to the two singlet states.

5. Conclusion

We have applied a GL-based perturbation theory along a range-separated adiabatic connection for the calculation of electronic excitation energies. Unlike the RS-based perturbation theory that we explored in a previous work, the GL-based perturbation theory keeps the ground-state density (and thus the ionization energy) constant at each order. Excitation energies up to first order in the perturbation have been calculated numerically for the helium and beryllium atoms and the hydrogen molecule without introducing any density-functional approximations.

In comparison with the RS-based perturbation theory, the GL-based perturbation theory gives much more accurate excitation energies for the Rydberg states of the helium atom but similar excitation energies for the valence states of the beryllium

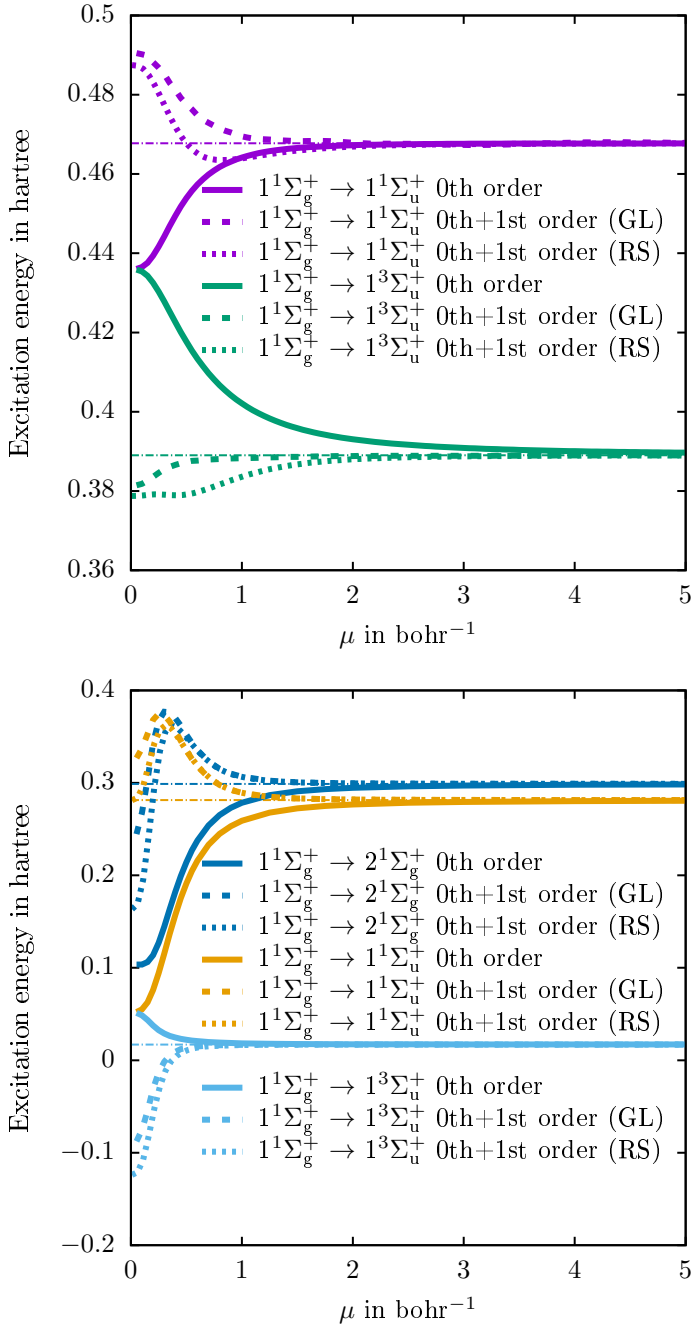

Figure 3. Zeroth-order excitation energies $\Delta\mathcal{E}_k^\mu = \mathcal{E}_k^\mu - \mathcal{E}_0^\mu$ (plain line) and zeroth+first order excitation energies $\Delta E_k^{\mu,(0+1)} = E_k^{\mu,(0+1)} - E_0^{\mu,(0+1)}$ (dashed line) in the GL-based perturbation theory for the hydrogen molecule at the equilibrium distance (top) and three times the equilibrium distance (bottom) as a function of μ. For comparison, the zeroth+first order excitation energies $\Delta E_{k,\mathrm{RS}}^{\mu,(0+1)} = E_{k,\mathrm{RS}}^{\mu,(0+1)} - E_{0,\mathrm{RS}}^{\mu,(0+1)}$ (dotted line) in the RS-based perturbation theory of Ref. [49] are also shown. The excitation energies of the physical system $\Delta E_k = E_k - E_0$ are plotted as thin horizontal dot-dashed lines.

atom and of the hydrogen molecule. This can be rationalized by observing that the Rydberg states are close to the ionization threshold and therefore sensitive to having the correct ionization energy.

This first-order GL-based perturbation theory works reasonably well for calculating

the first valence excitation energies of the hydrogen molecule at its equilibrium distance. However, results are less satisfactory for the valence excitation energies of the beryllium atom and stretched hydrogen molecule at small range-separation parameter. For such systems, with small HOMO-LUMO gaps, it may be necessary to go beyond single-reference first-order perturbation theory for small range-separated parameters.

One possible extension of this work would be to test density-functional approximations for the short-range exchange–correlation potential and approximations for the wave-function part of the calculation and perform more extensive benchmarking with a more practical implementation in a range-separated DFT code.

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