



Cite this: *Phys. Chem. Chem. Phys.*,
2024, 26, 6532

Signatures of s-wave scattering in bound electronic states

Robin E. Moorby,^{id} *^{ab} Valentina Parravicini,^{id} ^a Maristella Alessio^{id} ^a and
Thomas-C. Jagau^{id} *^a

We compute EOM-EA-CCSD and EOM-EA-CCSDT potential energy curves and one-electron properties of several anions at bond lengths close to where these states become unbound. We compare the anions of HCl and pyrrole, which are associated with s-wave scattering, with N₂ and H₂, which correspond to resonances. For HCl and pyrrole, we observe, on inclusion of diffuse basis functions, a pronounced bending effect in the anionic potential energy curves near the crossing points with their corresponding neutral molecules. Additionally, we observe that the Dyson orbital and second moment of the electron density become extremely large in this region; for HCl, the size of the latter becomes 5 orders of magnitude larger over a range of 5 pm. This behaviour is not observed in H₂ or N₂. Our work thus shows that bound state electronic-structure methods can distinguish between anions that turn into electronic resonances and those associated with s-wave scattering states.

Received 15th January 2024,
Accepted 30th January 2024

DOI: 10.1039/d4cp00181h

rsc.li/pccp

1 Introduction

The attachment of electrons to neutral molecules can lead to bound or unbound molecular anions. If the electron affinity of a molecule is positive, the attachment of the excess electron is permanent, and the anions can be treated theoretically with conventional Hermitian quantum-chemical methods. One can distinguish between valence anions, in which the additional electron resides in a rather compact orbital close to the molecule, and non-valence anions, where the additional electron resides in a diffuse orbital and is bound by either the molecular dipole moment, quadrupole moment, or dispersion effects.^{1–6} For the computational treatment of non-valence anions in particular, large basis sets are mandatory to capture the enormous extent of the wave function. Also, correlated wave function theory is often necessary to compute accurate electron affinities and, in some cases, an anion is not bound at all at the Hartree–Fock (HF) level of theory. Furthermore, it is notable that corrections to the energy from nuclear contributions can be largely different for the anionic and neutral molecules.^{1,7}

In cases where the anion is unbound at the equilibrium geometry of the neutral molecule, a non-zero angular momentum of the incoming electron can introduce a centrifugal barrier behind which the electron is temporarily trapped allowing for the formation of metastable electronic states, known as

shape resonances.^{1,8,9} The energy of electronic resonances places them within a continuum of elastically scattered states, meaning that they can decay *via* electron loss due to their coupling with the continuum.^{10–12} Such shape resonances are widespread, two well-characterised examples are the anions of H₂ and N₂. Besides shape resonances, there are Feshbach resonances which decay by a two electron process. A simple case of this type of state is the (σ_g)¹(σ_u)² excited state of H₂[–] at stretched bond lengths.^{13–15}

In the case of s-wave scattering, however, there is no centrifugal barrier. Temporary electron-attached states near the continuum threshold have been termed virtual states in s-wave scattering and, while also unbound, are distinct from resonances.^{8,9,16,17} Virtual states have been associated with anions in molecules such as CO₂,^{3,18–22} (CN)₂,²³ Fe(CO)₅,²⁴ HCl^{25–27} and pyrrole.²⁸

The asymptotic behaviour of unbound electronic states means that they cannot be treated with bound-state quantum chemical methods which fail to impose the correct boundary conditions. For electronic resonances, non-Hermitian quantum chemistry offers an elegant solution to this problem: the energies are calculated as complex-valued eigenvalues of a non-Hermitian Hamiltonian.^{10–12} Methods in this regard include complex scaling,^{29–31} where the coordinates of the Hamiltonian are rotated into the complex plane; complex basis functions, where the exponents of basis functions are rotated into the complex plane;³² and complex absorbing potentials,^{33,34} where a complex potential is applied to absorb the tails of the wave function. Alternatively, resonance positions can be determined using stabilisation and extrapolation

^a Department of Chemistry, KU Leuven, Celestijnenlaan 200F, B-3001 Leuven, Belgium. E-mail: robin.moorby@kuleuven.be, thomas.jagau@kuleuven.be

^b Department of Chemistry, University of Durham, South Road, Durham, DH1 3LE, UK



techniques.^{35–38} Analytical continuation of such methods is necessary to additionally calculate resonance decay widths.^{39,40}

These techniques can be integrated into bound-state quantum chemistry enabling a treatment of electronic resonances in analogy to that of bound states. However, their applicability to s-wave scattering and virtual states is questionable, which means that one must resort to more traditional methods based on scattering theory.⁹ The combination of the latter approaches with bound-state methods is still very challenging, even for diatomics such as HCl,^{26,27,41,42} which often necessitates the use of relatively low-level electronic-structure methods.

It is thus desirable to identify and characterise anions that correspond to virtual states and distinguish them from other anions that correspond to electronic resonances. To this end, we investigate in this work the anions of HCl and pyrrole both of which are relevant to s-wave scattering and contrast them with H_2^- and N_2^- as prototypical examples of resonances. While these anions all are unbound at the equilibrium structures of the respective neutral molecules, they become bound upon bond stretching. We show in this work that close to the point at which an anion becomes unbound, bound states turning into resonances and s-wave scattered states, respectively, exhibit different behaviour in terms of potential energy curves and one-electron properties.

2 Computational details

We conduct coupled-cluster singles and doubles (CCSD) calculations^{43,44} on the ground states of the neutral HCl, N_2 , H_2 , and pyrrole molecules and equation-of-motion electron-attachment CCSD (EOM-EA-CCSD) calculations^{45,46} to access the corresponding anions. We construct customised basis sets

from the aug-cc-pVTZ basis^{47–49} with additional diffuse functions obtained by recursively dividing the exponent of the most diffuse s and p shell by a factor of 2. A basis set labelled aug-cc-pVTZ+ n snp indicates that we have added n additional s and p shells to all atoms according to this even-tempered manner. In the largest basis sets used, aug-cc-pVTZ+15s15p, the smallest exponent, corresponding to an s function on hydrogen, is of the order of 10^{-7} . The molecular geometry of pyrrole is taken from ref. 50.

For all systems but pyrrole, we also carry out coupled-cluster singles, doubles and triples (CCSDT)^{51,52} and equation-of-motion electron-attachment CCSDT (EOM-EA-CCSDT) calculations.^{53,54} Our EOM-EA-CCSDT calculations make use of the continuum orbital method:⁵⁵ a basis function with an exponent of 10^{-10} , which effectively does not interact with the rest of the basis set, is included in the calculation and two electrons are placed in the resulting orbital when solving the restricted HF (RHF) equations. In a subsequent EOM excitation energy CCSDT (EOM-EE-CCSDT) calculation, one electron can then be excited from the continuum orbital into a target orbital allowing for the calculation of EOM-EA-CCSDT states using an EOM-EE-CCSDT implementation.

Additionally, for HCl^- and N_2^- , we compute the second moments of the electron density and the Dyson orbital of the anionic states at the EOM-EA-CCSD level. Dyson orbitals can be viewed as transition density matrices between neutral and electron-attached states and characterise electron attachment without invoking a mean-field approximation.^{56–58} The frozen-core approximation is used in all CC calculations. For HCl, we also compute the energy of the anion at the unrestricted HF (UHF) level of theory and by applying Koopmans' theorem to the virtual orbitals of the neutral molecule. All CCSD, EOM-EA-CCSD, and UHF calculations were performed using the Q-Chem software, version 6.0.2.⁵⁹ All CCSDT and EOM-EA-CCSDT

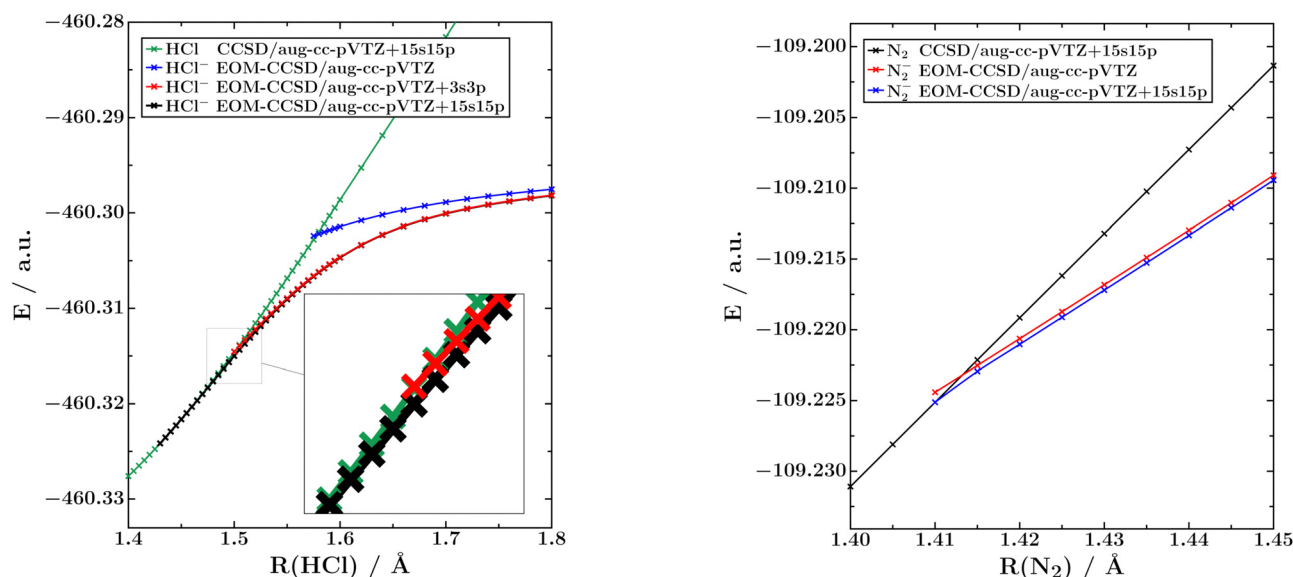


Fig. 1 Left: Potential energy curves of the ¹Σ⁺ ground state of HCl and the ²Σ⁺ ground state of HCl⁻ computed with CCSD and EOM-EA-CCSD, respectively, at 1.4 < R(HCl) < 1.8 Å. Right: Potential energy curves of the ¹Σ_g⁺ ground state of N₂ and the ²Π_g ground state of N₂⁻ computed with CCSD and EOM-EA-CCSD, respectively, at 1.4 < R(NN) < 1.45 Å.



calculations were performed using the CFOUR software, version 2.1.⁶⁰

3 Results

3.1 Potential energy curve of HCl^-

In the left panel of Fig. 1 we show potential energy curves of neutral HCl and of the $^2\Sigma^+$ ground state of HCl^- computed using CCSD and EOM-EA-CCSD, respectively, and three different basis sets: aug-cc-pVTZ, aug-cc-pVTZ+3s3p and aug-cc-pVTZ+15s15p. To illustrate the special character of these potential energy curves, we show in the right panel of the same figure the corresponding curves for neutral N_2 and the $^2\Pi_g$ ground state of N_2^- computed with the same methods and basis sets.

Fig. 1 demonstrates that the inclusion of diffuse shells in the basis set has a much bigger effect on HCl^- than on N_2^- . In the unmodified aug-cc-pVTZ basis, both anionic potential energy curves behave similarly, but upon the addition of diffuse shells the HCl^- curve bends downwards and crosses the curve of the neutral molecule at a much shorter H-Cl distance.

The crossing points between the neutral and anionic HCl potential energy curves computed with different methods and basis sets can be found in Table 1. It is seen that this point moves by 0.143 Å at the EOM-EA-CCSD level when going from the standard aug-cc-pVTZ basis to aug-cc-pVTZ+15s15p. In fact, we cannot assure that the value of 1.434 Å computed in the largest basis is converged with respect to basis-set size because the inclusion of 15 diffuse s and p shells represents a technical limit of our implementation. In contrast, the crossing point between the N_2 and N_2^- curves in the right panel of Fig. 1 is nearly invariant and only moves from 1.413 Å to 1.412 Å upon the inclusion of 15 diffuse s and p shells. We also note that the basis-set dependence of the HCl^- curve is only significant near the crossing point. At H-Cl bond lengths of 1.8 Å, the differences between the basis sets are negligible.

To confirm the validity of the EOM-EA-CCSD approximation for the HCl^- anion, we conducted EOM-EA-CCSDT calculations, which are documented in Table 1 as well. This illustrates that the crossing point moves to slightly shorter bond distances as compared to EOM-EA-CCSD but the bending effect remains similar in magnitude. However, due to technical limitations, we could not run EOM-EA-CCSDT calculations with more than 5 diffuse s and p shells. For N_2^- , differences between EOM-EA-CCSDT and EOM-EA-CCSD are very similar to those observed

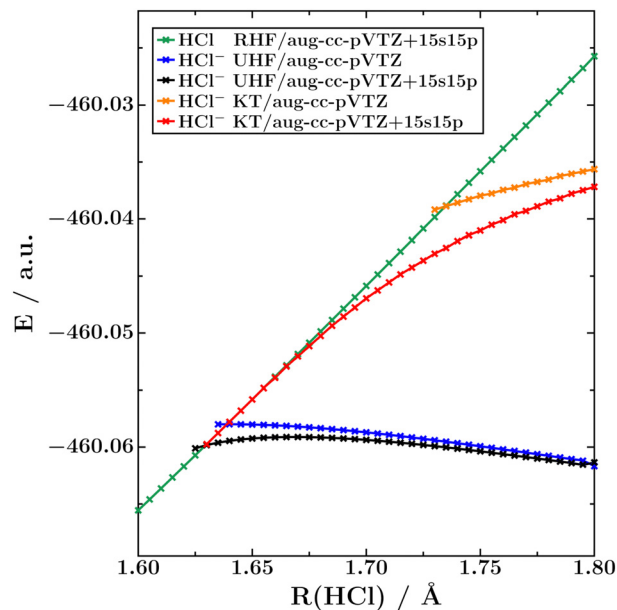


Fig. 2 Potential energy curves of the $^1\Sigma^+$ ground state of HCl and the $^2\Sigma^+$ ground state of HCl^- computed with HF theory and Koopmans' theorem at $1.6 < R(\text{HCl}) < 1.8$ Å.

for HCl^- , which corroborates that the bending feature observed for HCl^- is well captured by EOM-EA-CCSD and not related to an insufficient treatment of electron correlation.

To further investigate the nature of HCl^- , we also conducted UHF calculations on this anion and we also applied Koopmans' theorem to the lowest-lying virtual orbital of neutral HCl. The resulting potential energy curves are shown in Fig. 2, and the corresponding crossing points are available in Table 1. It is seen that UHF does not recover the bending effect and basis-set dependence visible in the EOM-CC potential energy curves from Fig. 1, while Koopmans' theorem does. At the HF level of theory, the inclusion of 15 diffuse s and p shells in the basis set moves the crossing point by only 0.009 Å, whereas that value amounts to 0.104 Å with Koopmans' theorem.

We add that there is a region of ca. 0.2 Å, from 1.63 Å to 1.434 Å, where HCl^- is only bound at correlated levels of theory. At these H-Cl distances, the energy of the anionic UHF potential curve lies above the RHF neutral potential curve, while the EOM-EA-CCSD anionic potential curve lies below the CCSD neutral potential curve. Also this behaviour is qualitatively different from N_2^- , which is bound already at shorter distances at the HF level of theory ($R(\text{NN}) \approx 1.33$ Å) than at the EOM-EA-CCSD level of theory ($R(\text{NN}) \approx 1.41$ Å).¹¹

We note that the dipole moment of HCl, computed with CCSD/aug-cc-pVTZ+15s15p, amounts to 1.48 D and 1.24 D at $R(\text{H-Cl}) = 1.8$ Å and 1.434 Å, respectively. This is substantially below the critical value needed to form a dipole-bound anion,¹ meaning that the polarity of HCl is not the critical difference to N_2 . However, the same bending effect shown for HCl^- in Fig. 1 and 2 has also been observed in EOM-EA-CCSD calculations along the bending coordinate of CO_2^- and connected to s-wave scattering.²⁰ A further interesting parallel can be drawn to

Table 1 Crossing points (Å) of the $^2\Sigma^+$ ground state of HCl^- with the $^1\Sigma^+$ ground state of HCl computed with EOM-EA-CCSD, EOM-EA-CCSDT, ΔHF, and Koopmans' theorem (KT) using basis sets with an increasing number of diffuse shells

Basis set	KT	ΔHF	CCSD	CCSDT
aug-cc-pVTZ	1.735	1.639	1.577	1.569
aug-cc-pVTZ+3s3p	1.682	1.630	1.502	1.498
aug-cc-pVTZ+5s5p	1.664	1.630	1.475	1.474
aug-cc-pVTZ+10s10p	1.642	1.630	1.445	
aug-cc-pVTZ+15s15p	1.631	1.630	1.434	



Table 2 Second moment of the electron density (\AA^2) of the $^2\Sigma^+$ ground state of HCl^- computed at various internuclear distances with EOM-EA-CCSD using basis sets with an increasing number of diffuse shells

Basis set	Crossing point	1.45 \AA	1.50 \AA	1.55 \AA	1.60 \AA	1.80 \AA
aug-cc-pVTZ	20.3 (1.577 \AA)				19.8	17.5
aug-cc-pVTZ+3s3p	71.4 (1.502 \AA)			48.6	31.4	18.0
aug-cc-pVTZ+5s5p	250.1 (1.475 \AA)		170.9	61.8	32.3	18.0
aug-cc-pVTZ+10s10p	6897.8 (1.445 \AA)	5507.7	255.2	62.5	32.4	18.6
aug-cc-pVTZ+15s15p	647 195.1 (1.434 \AA)	449 047.1	255.7	62.5	32.4	18.4

correlation-bound anions such as $\text{C}_6\text{F}_6^{-61}$ and C_{60}^{-62} also for these electronic states, a description based on a UHF wave function of the anion yields poor results, while EOM-EA-CC calculations based on an orbital manifold optimised for the neutral molecule perform much better.

3.2 Second moment of the electron density of HCl^-

To further characterise the HCl^- anion, we computed second moments of the electron density ($\langle r^2 \rangle$) with EOM-EA-CCSD and different basis sets. The results are reported in Table 2. At 1.8 \AA where HCl^- is in all basis sets bound in terms of Koopmans' theorem, $\langle r^2 \rangle$ is around 18 \AA². Increasing the number of diffuse shells has only a small impact on $\langle r^2 \rangle$ at this bond length. At 1.60 \AA, where HCl^- is bound at the EOM-EA-CCSD level but already unbound at the HF level, the basis-set dependence of $\langle r^2 \rangle$ is already more pronounced; 3 additional diffuse s and p shells on top of aug-cc-pVTZ lead to an increase by 50%. However, even more diffuse shells only become relevant at shorter distances: At 1.50 \AA, the value of $\langle r^2 \rangle$ has increased to 255 \AA² and 10 diffuse s and p shells are needed to capture it. At 1.434 \AA *i.e.*, the crossing point in our largest basis set (aug-cc-pVTZ+15s15p), we obtain a value of 647 000 \AA² for $\langle r^2 \rangle$, which corresponds to an average distance of *ca.* 800 \AA. Moreover, we cannot say whether this value is actually converged with respect to basis-set size.

The explosion of $\langle r^2 \rangle$ by a factor of more than 2×10^5 over a range of less than 0.2 \AA explains the pronounced basis-set dependence of the HCl^- potential energy curve in Fig. 1, but it is by no means typical of molecular anions. As a counter-example, we present in Table 3 $\langle r^2 \rangle$ values for N_2^- . For this anion, the second moment is nearly invariant with respect to basis-set changes and, in fact, decreases somewhat when approaching the crossing point.

3.3 Dyson orbital of HCl^-

The extremely diffuse nature of the EOM-EA-CCSD wave function of HCl^- near the crossing point with the neutral potential

Table 3 Second moment of the electron density (\AA^2) of the $^2\Pi_g$ ground state of N_2^- computed at various internuclear distances with EOM-EA-CCSD using basis sets with an increasing number of diffuse shells

Basis set	Crossing point	1.6 \AA	1.8 \AA
aug-cc-pVTZ	17.3	19.5	22.1
aug-cc-pVTZ+3s3p	17.7	19.5	22.1
aug-cc-pVTZ+5s5p	17.8	19.5	22.1
aug-cc-pVTZ+10s10p	17.9	19.5	22.1
aug-cc-pVTZ+15s15p	17.7	19.5	22.1

curve is also captured by the corresponding Dyson orbital, which is displayed in Fig. 3. Plotted at $R(\text{H-Cl}) = 1.434 \text{ \AA}$ and an isovalue of 5×10^{-5} (left panel of Fig. 3), the orbital resembles a σ^* orbital between hydrogen s and chlorine p atomic orbitals with a node in the centre of the bond. On reducing the isovalue to 3×10^{-5} (middle panel of Fig. 3), the character of the orbital changes and it becomes more akin to a σ orbital with a larger amplitude on the side of the hydrogen atom. At $R(\text{H-Cl}) = 1.8 \text{ \AA}$ (right panel of Fig. 3), the σ -character of the Dyson orbital is preserved but its spatial extent is considerably smaller consistent with the trend seen in $\langle r^2 \rangle$ in Table 2.

Similar to the second moment, the behaviour of the Dyson orbital of HCl^- is by no means typical of molecular anions as illustrated by Fig. 4, which shows the Dyson orbital of N_2^- . In contrast to HCl^- , the Dyson orbital of N_2^- remains compact and of π^* character independent of the isovalue and also when the N-N bond length is decreased from 1.8 \AA to 1.413 \AA, where the potential curves of N_2^- and N_2 cross. In fact, this orbital becomes slightly less diffuse when the bond becomes shorter. With non-Hermitian techniques, it is possible to evaluate a complex-valued Dyson orbital in the unbound region at $R(\text{NN}) < 1.413 \text{ \AA}$ as well;⁵⁷ the real part of such an orbital looks very similar to those displayed in Fig. 4.

The Dyson orbital of HCl^- shown in Fig. 3 is, however, similar to what one observes for correlation-bound anions.^{3,6,61,62} In C_6F_6^- , for example, the EOM-EA-CC natural orbital hosting the excess electron has a spatial extent much larger than the nuclear framework.⁶¹

3.4 Anions of H_2

There are two bound doublet states of H_2^- : a $^2\Sigma_u^+$ state, which is bound at bond lengths above *ca.* 1.6 \AA, and a $^2\Sigma_g^+$ state, which is bound at bond lengths above *ca.* 2.7 \AA. The configuration of the first state is $(\sigma_g)^2(\sigma_u)^1$, while that of the second state is $(\sigma_g)^1(\sigma_u)^2$. The first state turns into a shape resonance at shorter bond lengths, whereas the second state is first a Feshbach resonance and then becomes a shape resonance as soon as the $^3\Sigma_u^+$ state of H_2 is lower in energy.¹⁵ As H_2^- has only three electrons, both states can be treated exactly within a given basis set by means of EOM-EA-CCSDT calculations. Particularly relevant in the context of the present work is that the electron emitted in the decay of the $^2\Sigma_g^+$ state of H_2^- has to have zero angular momentum, meaning it represents an s-wave.

The potential curves of both anionic states computed with full EOM-EA-CC in different basis sets are shown in Fig. 5; the



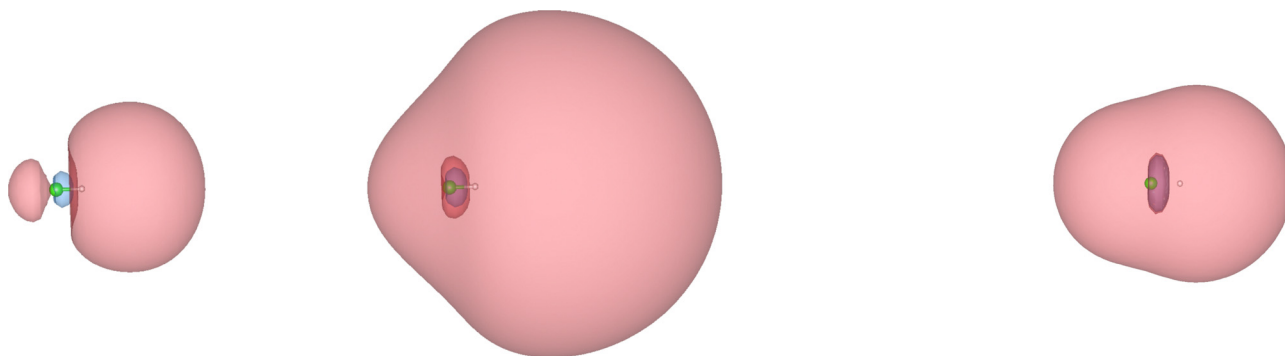


Fig. 3 Dyson orbital of the $2\Sigma^+$ ground state of HCl^- computed with EOM-EA-CCSD/aug-cc-pVTZ+15s15p. Left: $R(\text{HCl}) = 1.434 \text{ \AA}$, isovalue = 5×10^{-5} ; middle: $R(\text{HCl}) = 1.434 \text{ \AA}$, isovalue = 3×10^{-5} ; right: $R(\text{HCl}) = 1.8 \text{ \AA}$, isovalue = 1×10^{-3} .

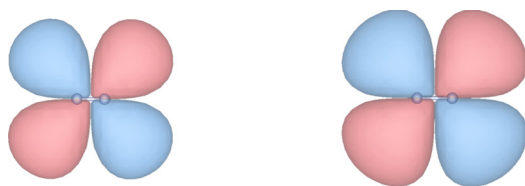


Fig. 4 Dyson orbital of the $2\Pi_g$ ground state of N_2^- computed with EOM-EA-CCSD/aug-cc-pVTZ+15s15p and plotted at an isovalue of 0.001 at internuclear distances of 1.413 Å (left) and 1.800 Å (right).

corresponding crossing points with the potential energy curve of neutral H_2 are summarised in Table 4. Fig. 5 shows that none of the two H_2^- states show the bending effect observed in HCl^- . Also, the position of the crossing point between the neutral and anionic potential curves depends much less on the basis set than in the case of HCl^- .

For the $2\Sigma_u^+$ state, the position of the crossing point is converged in the aug-cc-pVTZ+3s3p basis, where it differs by

Table 4 Crossing points (Å) of the $2\Sigma_u^+$ and $2\Sigma_g^+$ states of H_2^- with the $1\Sigma_g^+$ ground state of H_2 computed with EOM-EA-CCSD and EOM-EA-CCSDT using basis sets with an increasing number of diffuse shells. For EOM-EA-CCSD calculations on the ungerade state, we use the $3\Sigma_u^+$ state of H_2 as reference to enable a description as singly-excited state

Basis set	$2\Sigma_u^+$		$2\Sigma_g^+$	
	CCSD	CCSDT	CCSD	CCSDT
aug-cc-pVTZ	1.613	1.585	2.760	2.741
aug-cc-pVTZ+3s3p	1.594	1.570	2.672	2.641
aug-cc-pVTZ+5s5p	1.593		2.668	2.627
aug-cc-pVTZ+10s10p	1.593		2.668	
aug-cc-pVTZ+15s15p	1.593		2.668	

only 0.02 Å from the position in the aug-cc-pVTZ basis. The crossing point of the $2\Sigma_g^+$ state does vary more than that of the ungerade state when diffuse shells are added to the basis, but the position is converged within the aug-cc-pVTZ+5s5p basis and additional diffuse shells make no impact. We note that at

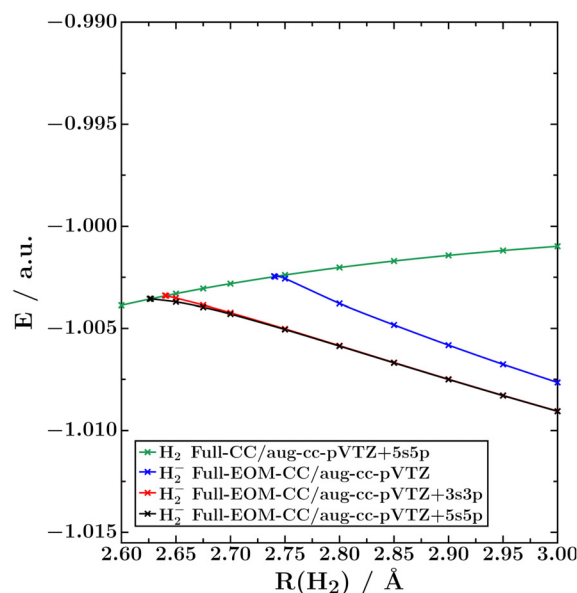
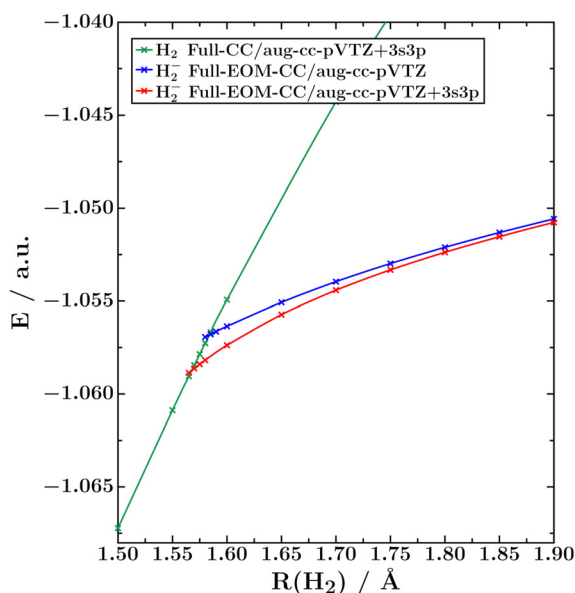


Fig. 5 Potential energy curves of the $2\Sigma_u^+$ (left) and $2\Sigma_g^+$ (right) states of H_2^- computed with full EOM-EA-CC. The $1\Sigma_g^+$ ground state of H_2 is also shown.



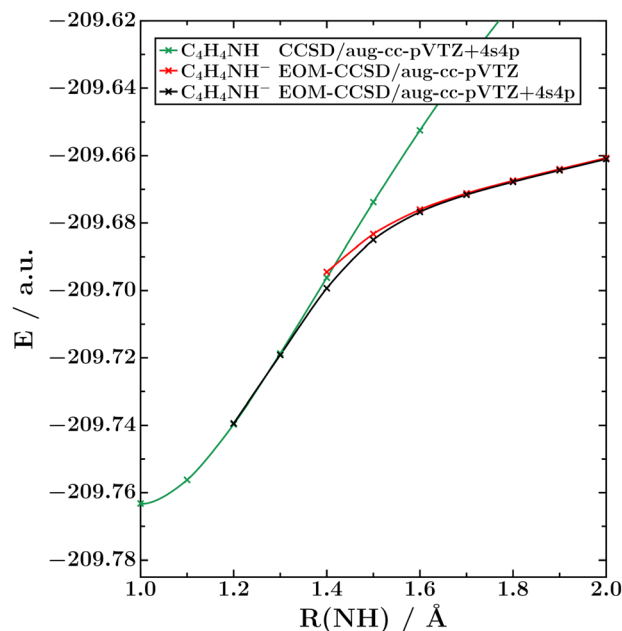


Fig. 6 Potential energy curves of the 1A_1 ground state of pyrrole and the 2A_1 state of the pyrrole anion computed with CCSD and EOM-EA-CCSD, respectively, as a function of N–H bond length at $1.0 < R(\text{NH}) < 2.0$ Å.

much longer bond distances of 4.0 and 5.0 Å the potential curves obtained in the aug-cc-pVTZ and aug-cc-pVTZ+5s5p basis sets continue to run parallel.

This indicates that the aug-cc-pVTZ basis is insufficient to describe the $^2\Sigma_g^+$ state at any bond length and the discrepancy in the position of the crossing points is not reflective of the bending effect observed in HCl^- , for which diffuse shells make very little impact at long bond distances. We finally note that the differences between EOM-EA-CCSD and EOM-EA-CCSDT are for both states of H_2^- similar to those observed for HCl^- and N_2^- .

3.5 Pyrrole anion

To demonstrate the relevance of our findings beyond diatomic molecules, we investigated the 2A_1 state of the pyrrole anion. Pyrrole does not support a bound anion at its equilibrium structure but has a rich electron-induced chemistry.^{28,50,63–65} While the nature of two π^* resonances is not debated, it is less clear if the low-lying totally symmetric anionic state, which is also present in pyrrole, should be interpreted as virtual state or rather as σ^* resonance.^{28,50,65} Independent of the nature of the latter state, there are indications that non-adiabatic transitions⁶⁶ between the π^* resonances and the totally symmetric state mediate dissociative electron attachment, which has implications for similar processes in more complex molecules.^{1,17}

For a comprehensive investigation of the pyrrole anion, the full-dimensional potential energy surface would need to be studied. By means of analytic gradient techniques,^{67–69} this is, in principle, possible also in the unbound regions as recent applications illustrated.⁵⁰ However, here we concentrate on the bound regions of the potential energy surface and furthermore limit ourselves to the N–H stretching coordinate.

As Fig. 6 illustrates, the bound part of the potential energy curve of the 2A_1 state of the pyrrole anion shows the same bending effect and basis-set dependence that we documented for HCl^- in Fig. 1 and that was previously observed in CO_2^- .²⁰ At the EOM-EA-CCSD level, the 2A_1 state is bound at $R(\text{NH}) > 1.2$ Å in the aug-cc-pVTZ+4s4p basis, but only at $R(\text{NH}) > 1.4$ Å in the unmodified aug-cc-pVTZ basis. Similar to HCl^- , the diffuse functions make no substantial impact at larger N–H distances of around 2 Å. This strongly suggests that the 2A_1 state turns into a virtual state in the unbound region and not a σ^* resonance.

4 Conclusions

We have investigated the bound anions of HCl, pyrrole, N_2 , and H_2 using the EOM-EA-CCSD and EOM-EA-CCSDT methods at molecular structures close to where they become unbound. The $^2\Sigma_g^+$ state of HCl^- and the 2A_1 state of pyrrole anion, both of which are connected to s-wave scattering, show a number of features that are not present in N_2^- and H_2^- : close to the crossing point with the parent neutral state, the spatial extent of the anionic wave function increases exponentially, which renders the potential energy curve extremely sensitive to diffuse functions in the basis set. For HCl^- , the inclusion of 15 diffuse s and p shells with exponents down to 10^{-7} on top of the aug-cc-pVTZ basis set yields a value of $647\,000$ Å² for the second moment of the electron density and moves the crossing point with the potential energy curve of neutral HCl by 0.143 Å compared to aug-cc-pVTZ. Likely, these values are not yet converged with respect to the size of the basis set. Also, they do not take into account effects beyond the Born–Oppenheimer approximation. In contrast, the $^2\Pi_g$ state of N_2^- and the $^2\Sigma_u^+$ state of H_2^- , which are associated with d-wave scattering and p-wave scattering, respectively, are not sensitive to diffuse basis functions as their spatial extent does not change by a lot along the potential energy curve. Interestingly, the totally-symmetric $^2\Sigma_g^+$ state of H_2^- , which is subject to a two-electron decay process past the crossing point, also does not show the features observed in HCl^- and the anion of pyrrole.

We note that the sensitivity of the potential energy curve towards diffuse basis functions was also observed for CO_2^- ,²⁰ another state associated with s-wave scattering. The unusual character of the HCl^- anion is also visible in the Dyson orbital, which, close to the crossing point, acquires an extremely diffuse character that is akin to correlation-bound anions of, for example, C_{60} or C_6F_6 , which have been related to s-wave scattering as well.^{3,61,62}

Importantly, the non-totally symmetric states of H_2^- and N_2^- , which turn into electronic shape resonances at shorter bond lengths, can be treated by means of non-Hermitian techniques such as complex basis functions or complex absorbing potentials in the unbound region.⁷⁰ The same applies to the totally-symmetric $^2\Sigma_g^+$ state of H_2^- , which turns into a Feshbach resonance.¹⁵

In contrast, the totally-symmetric anions of HCl, CO_2 , pyrrole, and similar molecules likely should not be considered as



electronic resonances in the unbound region. Instead, the term “virtual state” has been coined for them. Whether these virtual states are amenable to a treatment in terms of complex absorbing potentials or complex basis functions, or how to integrate them by other techniques into bound-state electronic-structure theory, is not clear at present and we hope that our work stimulates further research in this direction.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

R. E. M. acknowledges the European Union for mediating a bilateral exchange from the University of Durham to KU Leuven as a part of the Erasmus+ program. This work has been supported by a Marie Skłodowska-Curie Actions fellowship to M. A. (Grant Agreement No. 101062717). T.-C. J. gratefully acknowledges funding from the European Research Council (ERC) under the European Unions Horizon 2020 research and innovation program (Grant Agreement No. 851766) and the KU Leuven internal funds (Grant No. C14/22/083).

Notes and references

- 1 J. Simons, *J. Phys. Chem. A*, 2008, **112**, 6401–6511.
- 2 J. M. Herbert, *Rev. Comp. Chem.*, 2015, **28**, 391–517.
- 3 V. K. Voora, A. Kairalapova, T. Sommerfeld and K. D. Jordan, *J. Chem. Phys.*, 2017, **147**, 214114.
- 4 J. P. Rogers, C. S. Anstöter and J. R. R. Verlet, *Nat. Chem.*, 2018, **10**, 341–346.
- 5 J. Simons, *J. Phys. Chem. A*, 2023, **127**, 3940–3957.
- 6 G. P. Paran, C. Utku and T.-C. Jagau, *Phys. Chem. Chem. Phys.*, 2024, **26**, 1809–1818.
- 7 S. Gulania, T.-C. Jagau, A. Sanov and A. I. Krylov, *Phys. Chem. Chem. Phys.*, 2020, **22**, 5002–5010.
- 8 K. McVoy, *Nucl. Phys. A*, 1968, **115**, 481–494.
- 9 J. R. Taylor, *Scattering theory: the quantum theory of non-relativistic collisions*, John Wiley & Sons, Inc., Hoboken, NJ, 1972.
- 10 N. Moiseyev, *Non-Hermitian Quantum Mechanics*, Cambridge University Press, 2011.
- 11 T.-C. Jagau, K. B. Bravaya and A. I. Krylov, *Annu. Rev. Phys. Chem.*, 2017, **68**, 525–553.
- 12 T.-C. Jagau, *Chem. Commun.*, 2022, **58**, 5205–5224.
- 13 J. N. Bardsley and J. S. Cohen, *J. Phys. B: At., Mol. Opt. Phys.*, 1978, **11**, 3645.
- 14 D. T. Stibbe and J. Tennyson, *J. Phys. B: At., Mol. Opt. Phys.*, 1998, **31**, 815.
- 15 A. F. White, E. Epifanovsky, C. W. McCurdy and M. Head-Gordon, *J. Chem. Phys.*, 2017, **146**, 234107.
- 16 W. Domcke, *Phys. Rep.*, 1991, **208**, 97–188.
- 17 H. Hotop, M.-W. Ruf, M. Allan and I. Fabrikant, *Adv. At., Mol., Opt. Phys.*, 2003, **49**, 85–216.
- 18 M. A. Morrison, *Phys. Rev. A*, 1982, **25**, 1445–1449.
- 19 H. Estrada and W. Domcke, *J. Phys. B: At. Mol. Phys.*, 1985, **18**, 4469–4479.
- 20 T. Sommerfeld, *J. Phys. B: At., Mol. Opt. Phys.*, 2003, **36**, L127.
- 21 J. Dvořák, M. Ranković, K. Houfek, P. Nag, R. Čurík, J. Fedor and M. Čížek, *Phys. Rev. Lett.*, 2022, **129**, 013401.
- 22 J. Dvořák, K. Houfek and M. Čížek, *Phys. Rev. A*, 2022, **105**, 062821.
- 23 P. Nag, R. Čurík, M. Tarana, M. Polášek, M. Ehara, T. Sommerfeld and J. Fedor, *Phys. Chem. Chem. Phys.*, 2020, **22**, 23141–23147.
- 24 M. Allan, M. Lacko, P. Papp, Š. Matejčík, M. Zlatar, I. I. Fabrikant, J. Kočíšek and J. Fedor, *Phys. Chem. Chem. Phys.*, 2018, **20**, 11692–11701.
- 25 R. K. Nesbet, *J. Phys. B: At. Mol. Phys.*, 1977, **10**, L739.
- 26 M. Čížek, J. Horáček and W. Domcke, *Phys. Rev. A*, 1999, **60**, 2873–2881.
- 27 J. Fedor, C. Winstead, V. McKoy, M. Čížek, K. Houfek, P. Kolorenč and J. Horáček, *Phys. Rev. A: At., Mol., Opt. Phys.*, 2010, **81**, 042702.
- 28 T. P. Ragesh Kumar, P. Nag, M. Ranković, T. F. M. Luxford, J. Kočíšek, Z. Mašín and J. Fedor, Coupling of distant parts of pyrrole molecule through virtual and resonant states, *arXiv*, 2022, preprint, arXiv:2208.13539, DOI: [10.48550/arXiv.2208.13539](https://doi.org/10.48550/arXiv.2208.13539).
- 29 J. Aguilar and J. M. Combes, *Commun. Math. Phys.*, 1971, **22**, 269–279.
- 30 E. Balslev and J. M. Combes, *Commun. Math. Phys.*, 1971, **22**, 280–294.
- 31 B. Simon, *Commun. Math. Phys.*, 1972, **27**, 1–9.
- 32 C. W. McCurdy and T. N. Rescigno, *Phys. Rev. Lett.*, 1978, **41**, 1364–1368.
- 33 G. Jolicard and E. J. Austin, *Chem. Phys. Lett.*, 1985, **121**, 106–110.
- 34 U. V. Riss and H.-D. Meyer, *J. Phys. B: At., Mol. Opt. Phys.*, 1993, **26**, 4503–4535.
- 35 B. Nestmann and S. D. Peyerimhoff, *J. Phys. B: At. Mol. Phys.*, 1985, **18**, 615–626.
- 36 B. M. Nestmann and S. D. Peyerimhoff, *J. Phys. B: At. Mol. Phys.*, 1985, **18**, 4309–4319.
- 37 H. S. Taylor, in *Models, Interpretations, and Calculations Concerning Resonant Electron Scattering Processes in Atoms and Molecules*, John Wiley & Sons, Ltd, 1970, pp. 91–147.
- 38 A. U. Hazi and H. S. Taylor, *Phys. Rev. A*, 1970, **1**, 1109–1120.
- 39 J. Simons, *J. Chem. Phys.*, 1981, **75**, 2465–2467.
- 40 J. Simons, *J. Phys. Chem. A*, 2021, **125**, 7735–7749.
- 41 W. Domcke and C. Mündel, *J. Phys. B: At. Mol. Phys.*, 1985, **18**, 4491–4509.
- 42 M. Allan, M. Čížek, J. Horáček and W. Domcke, *J. Phys. B: At., Mol. Opt. Phys.*, 2000, **33**, L209–L213.
- 43 G. D. Purvis III and R. J. Bartlett, *J. Chem. Phys.*, 1982, **76**, 1910–1918.
- 44 I. Shavitt and R. J. Bartlett, *Many-Body Methods in Chemistry and Physics: MBPT and Coupled-Cluster Theory*, Cambridge University Press, Cambridge, UK, 2009.
- 45 J. F. Stanton and R. J. Bartlett, *J. Chem. Phys.*, 1993, **98**, 7029–7039.



- 46 M. Nooijen and R. J. Bartlett, *J. Chem. Phys.*, 1995, **102**, 3629–3647.
- 47 T. H. Dunning, *J. Chem. Phys.*, 1989, **90**, 1007–1023.
- 48 R. A. Kendall, T. H. Dunning and R. J. Harrison, *J. Chem. Phys.*, 1992, **96**, 6796–6806.
- 49 D. E. Woon and T. H. Dunning, *J. Chem. Phys.*, 1993, **98**, 1358–1371.
- 50 M. Mukherjee, T. P. Ragesh Kumar, M. Ranković, P. Nag, J. Fedor and A. I. Krylov, *J. Chem. Phys.*, 2022, **157**, 204305.
- 51 J. Noga and R. J. Bartlett, *J. Chem. Phys.*, 1987, **86**, 7041–7050.
- 52 J. Noga and R. J. Bartlett, *J. Chem. Phys.*, 1988, **89**, 3401.
- 53 S. A. Kucharski, M. Włoch, M. Musiał and R. J. Bartlett, *J. Chem. Phys.*, 2001, **115**, 8263–8266.
- 54 Y. J. Bomble, K. W. Sattelmeyer, J. F. Stanton and J. Gauss, *J. Chem. Phys.*, 2004, **121**, 5236–5240.
- 55 J. F. Stanton and J. Gauss, *J. Chem. Phys.*, 1999, **111**, 8785–8788.
- 56 C. Melania Oana and A. I. Krylov, *J. Chem. Phys.*, 2007, **127**, 234106.
- 57 T.-C. Jagau and A. I. Krylov, *J. Chem. Phys.*, 2016, **144**, 054113.
- 58 A. I. Krylov, *J. Chem. Phys.*, 2020, **153**, 080901.
- 59 E. Epifanovsky, A. T. B. Gilbert, X. Feng, J. Lee, Y. Mao, N. Mardirossian, P. Pokhilko, A. F. White, M. P. Coons, A. L. Dempwolff, Z. Gan, D. Hait, P. R. Horn, L. D. Jacobson, I. Kaliman, J. Kussmann, A. W. Lange, K. U. Lao, D. S. Levine, J. Liu, S. C. McKenzie, A. F. Morrison, K. D. Nanda, F. Plasser, D. R. Rehn, M. L. Vidal, Z.-Q. You, Y. Zhu, B. Alam, B. J. Albrecht, A. Aldossary, E. Alguire, J. H. Andersen, V. Athavale, D. Barton, K. Begam, A. Behn, N. Bellonzi, Y. A. Bernard, E. J. Berquist, H. G. A. Burton, A. Carreras, K. Carter-Fenk, R. Chakraborty, A. D. Chien, K. D. Closser, V. Cofer-Shabica, S. Dasgupta, M. de Wergifosse, J. Deng, M. Diedenhofen, H. Do, S. Ehlert, P.-T. Fang, S. Fatehi, Q. Feng, T. Friedhoff, J. Gayvert, Q. Ge, G. Gidofalvi, M. Goldey, J. Gomes, C. E. González-Espinoza, S. Gulania, A. O. Gunina, M. W. D. Hanson-Heine, P. H. P. Harbach, A. Hauser, M. F. Herbst, M. Hernández Vera, M. Hodecker, Z. C. Holden, S. Houck, X. Huang, K. Hui, B. C. Huynh, M. Ivanov, Á. Jász, H. Ji, H. Jiang, B. Kaduk, S. Kähler, K. Khistyayev, J. Kim, G. Kis, P. Klunzinger, Z. Koczor-Benda, J. H. Koh, D. Kosenkov, L. Koulias, T. Kowalczyk, C. M. Krauter, K. Kue, A. Kunitsa, T. Kus, I. Ladjánszki, A. Landau, K. V. Lawler, D. Lefrançois, S. Lehtola, R. R. Li, Y.-P. Li, J. Liang, M. Liebenthal, H.-H. Lin, Y.-S. Lin, F. Liu, K.-Y. Liu, M. Loipersberger, A. Luenser, A. Manjanath, P. Manohar, E. Mansoor, S. F. Manzer, S.-P. Mao, A. V. Marenich, T. Markovich, S. Mason, S. A. Maurer, P. F. McLaughlin, M. F. S. J. Menger, J.-M. Mewes, S. A. Mewes, P. Morgante, J. W. Mullinax, K. J. Oosterbaan, G. Paran, A. C. Paul, S. K. Paul, F. Pavošević, Z. Pei, S. Prager, E. I. Proynov, Á. Rák, E. Ramos-Cordoba, B. Rana, A. E. Rask, A. Rettig, R. M. Richard, F. Rob, E. Rossomme, T. Scheele, M. Scheurer, M. Schneider, N. Sergueev, S. M. Sharada, W. Skomorowski, D. W. Small, C. J. Stein, Y.-C. Su, E. J. Sundstrom, Z. Tao, J. Thirman, G. J. Tornai, T. Tsuchimochi, N. M. Tubman, S. P. Veccham, O. Vydrov, J. Wenzel, J. Witte, A. Yamada, K. Yao, S. Yeganeh, S. R. Yost, A. Zech, I. Y. Zhang, X. Zhang, Y. Zhang, D. Zuev, A. Aspuru-Guzik, A. T. Bell, N. A. Besley, K. B. Bravaya, B. R. Brooks, D. Casanova, J.-D. Chai, S. Coriani, C. J. Cramer, G. Cserey, A. E. DePrince III, R. A. DiStasio, Jr, A. Dreuw, B. D. Dunietz, T. R. Furlani, W. A. Goddard III, S. Hammes-Schiffer, T. Head-Gordon, W. J. Hehre, C.-P. Hsu, T.-C. Jagau, Y. Jung, A. Klamt, J. Kong, D. S. Lambrecht, W. Liang, N. J. Mayhall, C. W. McCurdy, J. B. Neaton, C. Ochsenfeld, J. A. Parkhill, R. Peverati, V. A. Rassolov, Y. Shao, L. V. Slipchenko, T. Stauch, R. P. Steele, J. E. Subotnik, A. J. W. Thom, A. Tkatchenko, D. G. Truhlar, T. Van Voorhis, T. A. Wesolowski, K. B. Whaley, H. L. Woodcock III, P. M. Zimmerman, S. Faraji, P. M. W. Gill, M. Head-Gordon, J. M. Herbert and A. I. Krylov, *J. Chem. Phys.*, 2021, **155**, 084801.
- 60 D. A. Matthews, L. Cheng, M. E. Harding, F. Lipparini, S. Stopkowicz, T.-C. Jagau, P. G. Szalay, J. Gauss and J. F. Stanton, *J. Chem. Phys.*, 2020, **152**, 214108.
- 61 V. K. Voora and K. D. Jordan, *J. Phys. Chem. A*, 2014, **118**, 7201–7205.
- 62 V. K. Voora, L. S. Cederbaum and K. D. Jordan, *J. Phys. Chem. Lett.*, 2013, **4**, 849–853.
- 63 A. Modelli and P. D. Burrow, *J. Phys. Chem. A*, 2004, **108**, 5721–5726.
- 64 E. M. de Oliveira, M. A. P. Lima, M. H. F. Bettega, S. d'A. Sanchez, R. F. da Costa and M. T. do N. Varela, *J. Chem. Phys.*, 2010, **132**, 204301.
- 65 S. A. Pshenichnyuk, I. I. Fabrikant, A. Modelli, S. Ptasinska and A. S. Komolov, *Phys. Rev. A*, 2019, **100**, 012708.
- 66 K. Chatterjee, Z. Koczor-Benda, X. Feng, A. I. Krylov and T.-C. Jagau, *J. Chem. Theory Comput.*, 2023, **19**, 5821–5834.
- 67 Z. Benda and T.-C. Jagau, *J. Chem. Phys.*, 2017, **146**, 031101.
- 68 Z. Benda, K. Rickmeyer and T.-C. Jagau, *J. Chem. Theory Comput.*, 2018, **14**, 3468–3478.
- 69 Z. Benda and T.-C. Jagau, *J. Chem. Theory Comput.*, 2018, **14**, 4216–4223.
- 70 T.-C. Jagau and A. I. Krylov, *J. Phys. Chem. Lett.*, 2014, **5**, 3078–3085.

