

DENSITY CUMULANT THEORY FOR GROUND AND EXCITED ELECTRONIC STATES

by

ANDREAS VICTOR COPAN

(Under the Direction of Henry F. Schaefer III)

ABSTRACT

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INDEX WORDS: Cool stuff, Deep learning, Big data, Crowdsourcing, Computer vision, Computer science

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FOR GROUND AND EXCITED ELECTRONIC STATES

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ANDREAS VICTOR COPAN

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A Dissertation Submitted to the Graduate Faculty
of The University of Georgia in Partial Fulfillment
of the
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DOCTOR OF PHILOSOPHY

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In memory of Valery Andreiyevich and Valtraut Kirsch Copan



Acknowledgments

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Contents

Acknowledgments	v
1 Introduction	1
1.1 Stuff	1
2 Ground-State Density Cumulant Theory:	
Thermochemical and Kinetic Benchmark Calculations	3
2.1 Abstract	3
2.2 Introduction	4
2.3 Overview of DCFT	7
2.4 Computational Details	9
2.5 Results and Discussion	9
2.6 Conclusions	33
3 Linear-Response Density Cumulant Theory for Excited States:	
First Implementation and Benchmark Calculations	36
3.1 Abstract	36
3.2 Introduction	37

3.3	Theory	40
3.4	Computational Details	49
4	Linear-Response Density Cumulant Theory for Excited States:	
	Better Algorithms, Bigger Systems	51
5	Conclusion	55
	Appendices	58
A	Extended Results	58
	Bibliography	61

Chapter 1

Introduction

Here is the introduction. My dissertation introduces *widgets*. If I wish to cite someone, I could refer to them in text as in Pedersen, Fernández, and Koch¹, or parenthetically¹. I can also cite just the year 2001 or mention the author, Pedersen, Fernández, and Koch.

This dissertation template is handy. I like it and use it every time I write a dissertation.

— Abraham Lincoln 2001

1.1 Stuff

Here is a section on stuff.

1.1.1 Artificial Neural Networks

See Figure 1.1 for a neural network.

Cool.

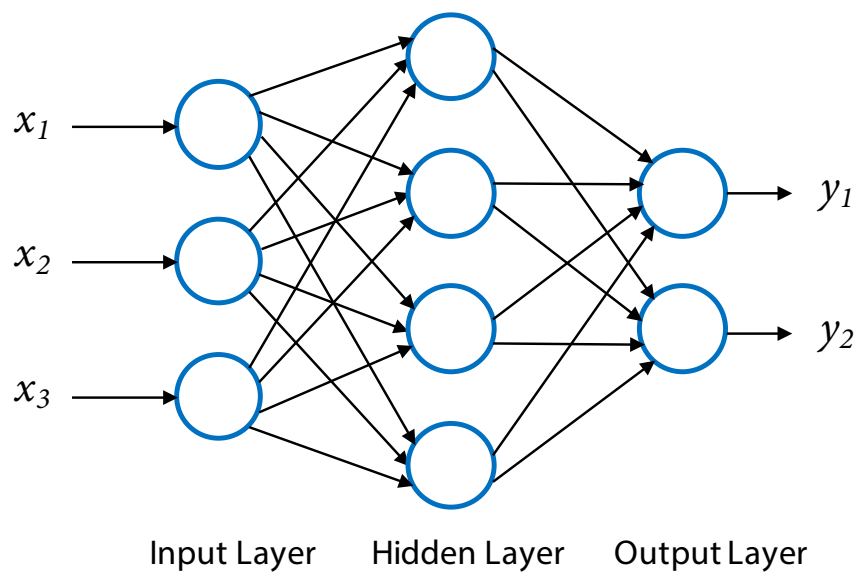


Figure 1.1: An example neural network with two final outputs. Notice how each neuron in one layer connects to each neuron in the following layer. This is called *fully connected*.

Chapter 2

Ground-State Density Cumulant Theory: Thermochemical and Kinetic Benchmark Calculations*

2.1 Abstract

We present an extensive benchmark study of density cumulant functional theory (DCFT) for thermochemistry and kinetics of closed- and open-shell molecules. The performance of DCFT methods (DC-06, DC-12, ODC-06, and ODC-12) is compared to that of coupled-electron pair methods (CEPA₀ and OCEPA₀) and coupled-cluster theory (CCSD and CCSD(T)) for the description of noncovalent interactions (A24 database), barrier heights of hydrogen-transfer reactions (HTBH38), radical stabilization energies (RSE30), adiabatic ionization energies (AIE), and covalent bond stretching in diatomic molecules. Our results indicate that out of four DCFT methods the ODC-12 method is the most reliable and accurate DCFT formulation to date. Compared to CCSD, ODC-12 shows superior results for all benchmark tests employed in our study. With respect to coupled-pair

*A. V. Copan, A. Yu. Sokolov, and H. F. Schaefer., J. Chem. Theory Comput. **10**, 2389 (2014). Adapted with permission of the American Chemical Society.

theories, ODC-12 outperforms CEPA₀, and shows similar accuracy to the orbital-optimized CEPA₀ variant (OCEPA₀) for systems at equilibrium geometries. For covalent bond stretching, ODC-12 is found to be more reliable than OCEPA₀. For the RSE30 and AIE datasets, ODC-12 shows competitive performance with CCSD(T). In addition to benchmark results, we report new reference values for the RSE30 dataset computed using coupled cluster theory with up to perturbative quadruple excitations.

2.2 Introduction

Recent developments in *ab initio* quantum chemistry have resulted in a variety of computational models for studying molecules. Apart from concerns about efficiency and accuracy, several concepts have evolved as criteria for judging the merits of a particular method. Energy-based criteria typically define an “ideal” approximation as one yielding correlation energies that are size-consistent, extensive², well-defined (giving continuous, unique potential surfaces), and variational.³ While it has been argued that the practical benefits of variationality are rather limited,⁴ the efficiency of gradient computations, at least, is improved by formulating a theory in terms of a Hermitian and stationary energy functional.⁵ With respect to scope and stability, methods that show consistent performance for open-shell systems, strongly correlated states, and non-equilibrium geometries are particularly valuable.⁴

The incorrect scaling of truncated configuration interaction (CI) energies with system size has inspired the development of size-extensive alternatives. Among

the earliest formulations, the coupled electron pair approximations (CEPAs)^{6–10} attracted much attention in 1970s,^{11–15} offering rigorous extensivity and size-consistency while retaining much of the linearity¹⁶ of CI in their equations. CEPA methods, however, have been shown to rapidly deteriorate as the molecular geometry deviates from equilibrium¹⁶ and yield energies that vary under the rotation of the occupied orbitals.⁹ Partly in light of such defects, CEPA has been largely displaced by coupled-cluster (CC) theory.^{4;17–23} In addition to size-extensivity, CC offers orbital invariance and improved stability for non-equilibrium structures¹⁶, but has a non-Hermitian energy functional and non-linear equations which are not readily amenable to parallel implementation. Although neither class of methods is strictly variational, VCEPA (variational CEPA) has been shown to be effectively equivalent to its non-variational counterpart.²⁴ Various other modifications to resolve the deficiencies of traditional CEPA have been explored, including self-consistent size-consistent CI,^{25;26} orbital-invariant CEPA,^{27;28} and orbital-optimized CEPA formulations.^{29–32} Recently, the CEPA methods have been revived by Neese and co-workers^{24;33;34} who developed the local pair-natural-orbital CEPA (LPNO-CEPA) methods and have implemented them for massively parallel computer architectures.

It has recently been demonstrated^{35–38} that CEPA methods naturally arise in the context of theories that obtain the molecular energies from density cumulants, the connected and extensive components of the reduced density matrices (RDMs).^{39–44} The advantage of cumulant-based theories is that, unlike their RDM-based counterparts,^{45–47} they are naturally size-extensive and size-consistent.^{42;48}

We have recently achieved the first implementation^{49;50} of density cumulant functional theory (DCFT), proposed by Kutzelnigg in 2006.³⁵ In DCFT, the molecular energy is obtained in terms of a mean-field one-particle RDM and the two-particle density cumulant, constrained to be at least approximately N -representable (*i.e.* to correspond to a physical N -electron wavefunction). Like traditional CC theory, DCFT is size-extensive and orbital-invariant, but it has the additional advantage of a stationary and Hermitian energy functional, which simplifies the computation of molecular properties. In the original DCFT formulation (DC-06)^{35;49;50} N -representability conditions derived from second-order Møller-Plesset perturbation theory (MPPT) were used,⁵¹ yielding equations similar to those of the simplest CEPA model (CEPA₀),^{8;10} but including higher-order terms in the description of one-particle correlation effects. Using the same set of conditions, we have developed new formulations of DCFT that take advantage of an improved description of the one-particle density matrix (DC-12)⁵² and full orbital optimization (ODC-06 and ODC-12 methods).⁵³

Our previous studies^{49;50;52;53} demonstrated for a limited set of systems that the DC-06, DC-12, ODC-06 and ODC-12 methods generally yield molecular energies and properties competitive with those obtained by CCSD and CCSD(T), but may exhibit unstable performance due to imbalances in the description of electron correlation. Herein, we present an extensive benchmark of the DCFT methods with respect to thermochemical and kinetic molecular properties, including noncovalent interactions, barrier heights in hydrogen-transfer reactions, radical stabilization energies, and adiabatic ionization energies for challenging electron-

dense systems. We conclude our benchmark study by testing the performance of DCFT for covalent bond stretching in diatomic molecules.

2.3 Overview of DCFT

In this section a short overview of DCFT is presented. For details on the theory the reader is referred to our earlier publications.^{49;52;53} In the RDM methods⁵⁴ the exact molecular energy is expressed as a functional of the one- and two-particle reduced density matrices, γ_1 and γ_2 (1-RDM and 2-RDM):

$$E = h_p^q \gamma_q^p + \frac{1}{2} g_{pq}^{rs} \gamma_{rs}^{pq}, \quad [\gamma_1]_q^p \equiv \gamma_q^p, \quad [\gamma_2]_{rs}^{pq} \equiv \gamma_{rs}^{pq}. \quad (2.1)$$

In eq. (2.1), h_p^q and g_{pq}^{rs} are the usual one- and two-electron integrals in the orthonormal spin-orbital basis $\{\psi_p\}$ and summation over the repeated indices is implied. Expressing γ_1 through γ_2 via the partial trace relation $\sum_r \gamma_{qr}^{pr} = (N-1)\gamma_q^p$, the energy functional (2.1) can be minimized by varying γ_2 subject to N -representability constraints. This is the essence of the variational 2-RDM approach.⁵⁴

In DCFT, some of the challenges of the 2-RDM approach are circumvented by expanding γ_2 in terms of its irreducible components – the 1-RDM and the two-particle cumulant (denoted by λ_2):

$$\gamma_{rs}^{pq} = \gamma_r^p \gamma_s^q - \gamma_r^q \gamma_s^p + \lambda_{rs}^{pq}. \quad (2.2)$$

In eq. (2.2), λ_2 describes the correlated part of γ_2 that cannot be expressed via γ_1 . The cumulant also determines the correlation contribution to γ_1 , allowing

the 1-RDM to be decomposed as the sum of an idempotent 1-RDM ($\boldsymbol{\kappa}$) and a correlation correction ($\boldsymbol{\tau}$):

$$\boldsymbol{\gamma}_1 = \boldsymbol{\kappa} + \boldsymbol{\tau}. \quad (2.3)$$

The correlation component $\boldsymbol{\tau}$ is fully specified by $\boldsymbol{\lambda}_2$, whereas $\boldsymbol{\kappa}$ is independent of $\boldsymbol{\lambda}_2$. Equations (2.2) and (2.3) allow us to write an equivalent energy expression with $\boldsymbol{\kappa}$ and $\boldsymbol{\lambda}_2$ as independent functional parameters:

$$\begin{aligned} E[\boldsymbol{\kappa}, \boldsymbol{\lambda}_2] &= \frac{1}{2}(h_p^q + f_p^q)(\kappa_q^p + \tau_q^p) + \frac{1}{4}\bar{g}_{pq}^{rs}\lambda_{pq}^{rs}, \\ f_p^q &= h_p^q + \bar{g}_{pr}^{qs}(\kappa_s^r + \tau_s^r), \quad \bar{g}_{rs}^{pq} = g_{rs}^{pq} - g_{rs}^{qp}. \end{aligned} \quad (2.4)$$

Here, the generalized Fock operator \boldsymbol{f} differs from that of Hartree-Fock theory by the presence of an external potential $\bar{g}_{pr}^{qs}\tau_s^r$ due to electron correlation.³⁵

To date, all DCFT formulations make the energy (2.4) stationary with respect to variations of $\boldsymbol{\lambda}_2$, subject to cumulant N -representability constraints derived from second-order Møller-Plesset perturbation theory (MPPT).⁵¹ To account for orbital relaxation effects, the two earliest DCFT methods, DC-06^{35;49;50} and DC-12⁵², determined the orbitals by diagonalizing the generalized Fock operator \boldsymbol{f} defined in eq. (2.4). These two methods differ in their description of 1-RDM N -representability. Whereas DC-06 employs an approximate expression for $\boldsymbol{\tau}$ in terms of $\boldsymbol{\lambda}_2$, DC-12 uses the exact relationship. Recently, we proposed orbital-optimized variants of DC-06 and DC-12 (ODC-06 and ODC-12),⁵³ which fully account for orbital relaxation effects.

2.4 Computational Details

All computations were performed using the Psi4 package.⁵⁵ The results were benchmarked against coupled cluster theory with single and double excitations (CCSD)^{21–23}, CCSD with perturbative triple excitations [CCSD(T)],^{56;57} coupled electron pair approximation zero (CEPA₀),^{8;10} and the orbital-optimized variant of CEPA₀ (OCEPA₀)³⁰. All electrons were correlated in all computations. The cc-pCVXZ^{58;59} and aug-cc-pVXZ⁶⁰ basis sets (X = T, Q) were used (see text for details). Noncovalent interaction energies, hydrogen-transfer barrier heights, and radical stabilization energies were computed using geometries from the A24⁶¹, HTBH38⁶², and RSE30⁶³ benchmark databases, respectively, available in Psi4. Adiabatic ionization energies were computed from neutral and cation geometries optimized at each level of theory, with added harmonic zero-point vibrational energy corrections. Harmonic frequencies were computed by numerical differentiation of analytic energy gradients. Single-point energies were converged to $10^{-8} E_h$, while the root mean square of the energy gradient was converged to $10^{-6} E_h/a_0$ for geometry optimizations.

2.5 Results and Discussion

2.5.1 Noncovalent Interactions

We begin by testing the accuracy of DCFT methods for the description of noncovalent interactions in 24 closed-shell molecular dimers, which are listed in table 2.1. These molecular complexes comprise the A24 dataset⁶¹ developed by Řezáč and

Table 2.1: Errors in interaction energies (kcal mol⁻¹) for 24 noncovalently bound molecular dimers comprising the A24 database⁶¹ computed using seven methods with the aug-cc-pVTZ basis set. The errors are relative to CCSD(T) reference values (kcal mol⁻¹) shown in the rightmost column. For each method the mean absolute deviations from CCSD(T) (Δ_{MAE} , kcal mol⁻¹) and the standard deviations from the mean signed error (Δ_{SD} , kcal mol⁻¹) are also shown.

Complex (Sym.)	ΔCEPA_0	$\Delta\text{DC-06}$	$\Delta\text{DC-12}$	ΔCCSD	ΔOCEPA_0	$\Delta\text{ODC-06}$	$\Delta\text{ODC-12}$	CCSD(T)
H ₂ O...NH ₃ (C _s)	0.26	0.24	0.22	0.36	0.19	0.20	0.18	-7.18
H ₂ O...H ₂ O (C _s)	0.19	0.18	0.16	0.25	0.13	0.14	0.12	-5.71
HCN...HCN (C _s)	0.21	0.27	0.16	0.15	0.18	0.26	0.14	-7.12
HF...HF (C _s)	0.14	0.13	0.11	0.16	0.08	0.09	0.07	-5.20
NH ₃ ...NH ₃ (C _{2h})	0.15	0.13	0.14	0.26	0.12	0.12	0.12	-3.43
HF...CH ₄ (C _{3v})	0.17	0.16	0.20	0.23	0.12	0.12	0.16	-2.30
NH ₃ ...CH ₄ (C _{3v})	0.07	0.05	0.05	0.13	0.05	0.05	0.04	-1.08
H ₂ O...CH ₄ (C _s)	0.06	0.05	0.04	0.11	0.05	0.05	0.04	-1.03
CH ₂ O...CH ₂ O (C _s)	0.89	0.99	0.65	0.46	0.62	0.87	0.46	-5.23
H ₂ O...C ₂ H ₄ (C _s)	0.15	0.16	0.15	0.31	0.20	0.26	0.21	-3.33
CH ₂ O...C ₂ H ₄ (C _s)	0.21	0.18	0.14	0.27	0.19	0.24	0.16	-2.24
HCCH...HCCH (C _{2v})	0.07	0.05	0.05	0.20	0.10	0.12	0.10	-2.57

Complex (Sym.)	ΔCEPA_0	$\Delta\text{DC-06}$	$\Delta\text{DC-12}$	ΔCCSD	ΔOCEPA_0	$\Delta\text{ODC-06}$	$\Delta\text{ODC-12}$	CCSD(T)
$\text{NH}_3\cdots\text{C}_2\text{H}_4$ (C_s)	0.09	0.06	0.08	0.24	0.12	0.15	0.13	-2.07
$\text{C}_2\text{H}_4\cdots\text{C}_2\text{H}_4$ (C_{2v})	0.10	0.02	0.07	0.33	0.14	0.13	0.15	-1.81
$\text{CH}_4\cdots\text{C}_2\text{H}_4$ (C_s)	0.02	-0.02	0.01	0.14	0.05	0.04	0.06	-0.92
$\text{BH}_3\cdots\text{CH}_4$ (C_s)	0.23	0.18	0.24	0.37	0.18	0.16	0.22	-2.52
$\text{CH}_4\cdots\text{C}_2\text{H}_4$ (C_s)	0.13	0.09	0.13	0.23	0.10	0.09	0.09	-1.37
$\text{CH}_4\cdots\text{C}_2\text{H}_6$ (C_s)	0.09	0.06	0.09	0.17	0.07	0.06	0.09	-1.14
$\text{CH}_4\cdots\text{CH}_4$ (D_{3d})	0.08	0.06	0.08	0.14	0.06	0.05	0.08	-0.93
$\text{Ar}\cdots\text{CH}_4$ (C_{3v})	0.07	0.05	0.07	0.10	0.05	0.05	0.06	-0.78
$\text{Ar}\cdots\text{C}_2\text{H}_4$ (C_{2v})	0.03	-0.01	0.02	0.11	0.05	0.03	0.05	-0.63
$\text{C}_2\text{H}_4\cdots\text{HCCH}$ (C_{2v})	-0.02	-0.19	-0.01	0.38	0.07	-0.06	0.11	0.43
$\text{C}_2\text{H}_4\cdots\text{C}_2\text{H}_4$ (D_{2h})	-0.05	-0.30	-0.03	0.43	0.04	-0.16	0.11	0.41
$\text{HCCH}\cdots\text{HCCH}$ (D_{2h})	0.01	-0.09	0.02	0.34	0.10	0.02	0.12	0.91
$\Delta_{\text{MAE}}:$	0.14	0.16	0.12	0.25	0.13	0.15	0.13	
$\Delta_{\text{SD}}:$	0.18	0.23	0.13	0.11	0.12	0.18	0.09	

Hobza to include a variety of noncovalent interactions, including hydrogen bonding and π - π stacking. Although Řezáč and Hobza reported the interaction energies at the CCSD(T) complete basis set (CBS) limit, we use CCSD(T)/aug-cc-pVTZ energies as reference values in order to effectively exclude basis-set incompleteness error from the comparison.

Figure 2.1 depicts mean absolute error (Δ_{MAE}) relative to CCSD(T) in the binding energies of CEPA₀, OCEPA₀, CCSD, and the four DCFT methods (DC-06, DC-12, ODC-06, and ODC-12), as well as the root mean square deviation from the average signed error (Δ_{SD}). All methods but CCSD give similar Δ_{MAE} values (0.14 ± 0.02 kcal mol⁻¹), and a comparison between CEPA₀, DC-06, and DC-12 and their orbital-optimized variants (OCEPA₀, ODC-06, and ODC-12) shows negligible 0.01 kcal mol⁻¹ differences in each case. CCSD gives a significantly larger Δ_{MAE} (0.25 kcal mol⁻¹) than the other methods, exceeding the DC-12 Δ_{MAE} by a factor of two (0.12 kcal mol⁻¹). The Δ_{SD} values are much more sensitive to the choice of method than the Δ_{MAE} values, and are noticeably affected by orbital optimization. ODC-12 gives the smallest standard deviation ($\Delta_{\text{SD}} = 0.09$ kcal mol⁻¹), while the largest Δ_{SD} value was found for DC-06 (0.23 kcal mol⁻¹). The OCEPA₀, ODC-06, and ODC-12 methods ($\Delta_{\text{SD}} = 0.12, 0.18, \text{ and } 0.09$ kcal mol⁻¹, respectively) exhibit much more consistent performance than their non-orbital-optimized analogues, with Δ_{SD} smaller by 0.05 ± 0.01 kcal mol⁻¹ in each case. CCSD also exhibits a relatively small Δ_{SD} value (0.11 kcal mol⁻¹), possibly due to its inclusion of single excitations which partly account for orbital relaxation.

Errors in interaction energy and CCSD(T) reference values for each molecular

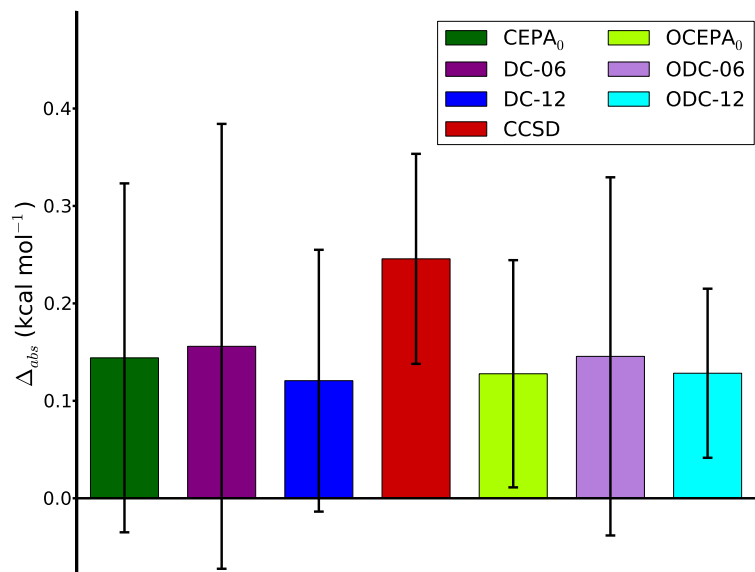


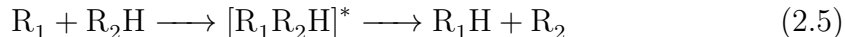
Figure 2.1: Mean absolute deviations (Δ_{MAE} , kcal mol⁻¹) and the standard deviations from the mean signed error (Δ_{SD} , kcal mol⁻¹) of the interaction energies for 24 noncovalently bound molecular dimers (A24 database) computed using seven methods with the aug-cc-pVTZ basis set. The errors are relative to CCSD(T)/aug-cc-pVTZ reference values. The Δ_{MAE} value is represented as a height of each colored box, while the Δ_{SD} value is depicted as a radius of the black vertical bar. See table 2.1 for data on individual database members.

complex are shown in table 2.1. The largest deviations from CCSD(T) were obtained for the formaldehyde dimer ($\text{CH}_2\text{O} \cdots \text{CH}_2\text{O}$, complex 9 in table 2.1), for which DC-06, CEPA₀, and OCEPA₀ yield errors of 0.99, 0.89, and 0.87 kcal mol⁻¹, respectively. For this system, the best performance is shown by CCSD and ODC-12, both of which give an error of 0.46 kcal mol⁻¹. For systems with π -stacking

interactions (complexes 22-24 in table 2.1), CCSD shows large errors (0.38, 0.43, 0.34 kcal mol⁻¹) relative to the magnitude of the interaction energy (0.43, 0.41, 0.91 kcal mol⁻¹, respectively). Here CEPA₀, DC-12, and their orbital-optimized variants offer much better agreement with CCSD(T), with errors ranging from 0.01 to 0.15 kcal mol⁻¹.

2.5.2 Hydrogen-Transfer Reaction Barrier Heights

We continue by assessing the performance of DCFT methods in predicting barrier heights for 18 hydrogen-transfer reactions from the HTBH38 database:⁶²



These reactions[†] involve molecules (R_1 and R_2) and transition states ($[R_1R_2H]^*$) with open-shell character, making their properties more sensitive to electron correlation effects. We employ barrier heights computed at the CCSD(T)/aug-cc-pVTZ level of theory as our reference rather than the values provided by Lynch⁶⁴ in order to effectively exclude basis-set incompleteness effects. We also omit the DC-06 and ODC-06 methods, which encounter frequent convergence problems due to the poor description of N -representability (see Supporting Information for incomplete DC-06 results).

Mean absolute deviations (Δ_{MAE}) and standard deviations (Δ_{SD}) for the hydrogen-transfer barrier heights are presented in table 2.2 and plotted in fig. 2.2. The

[†]Reaction 19 in HTBH38, the cis-trans isomerization of piperylene, is omitted in the present study.

Table 2.2: Errors in barrier heights (kcal mol⁻¹) for 18 hydrogen-transfer reactions ($R_1 + R_2H \rightarrow R_1H + R_2$) comprising the HTBH38 database⁶⁴ computed using five methods with the aug-cc-pVTZ basis set. The errors are relative to CCSD(T) reference values (kcal mol⁻¹) shown in the rightmost column. Each reaction includes barrier heights in the forward ($R_1 + R_2H \rightarrow [R_1R_2H]^*$) and reverse ($[R_1R_2H]^* \leftarrow R_1H + R_2$) directions, respectively, except in the case of $R_1 = R_2 = H$ where they are the same. The mean absolute (Δ_{MAE} , kcal mol⁻¹) and the mean percent (Δ_{rel} , %) errors with respect to CCSD(T), as well as the standard deviations from the mean signed error (Δ_{SD} , kcal mol⁻¹) are also shown.

	Reaction Barrier	$\Delta CEPA_0$	$\Delta DC-12$	$\Delta CCSD$	$\Delta OCEPA_0$	$\Delta ODC-12$	CCSD(T)
1	$H + HCl \rightarrow [HHCl]^*$	0.74	0.49	0.09	-0.41	-0.28	5.22
2	$OH + H_2 \rightarrow [OHH_2]^*$	3.77	3.38	1.82	0.88	1.24	4.99
3	$CH_3 + H_2 \rightarrow [CH_3H_2]^*$	1.60	1.46	1.37	0.46	0.70	11.29
4	$OH + CH_4 \rightarrow [OHCH_4]^*$	4.26	3.85	2.61	1.22	1.65	5.64
5	$H + H_2 \rightarrow [HH_2]^*$	0.80	0.69	0.30	-0.27	-0.05	9.77
6	$OH + NH_3 \rightarrow [OHNH_3]^*$	6.02	5.25	3.54	1.18	1.82	3.17
7	$HCl + CH_3 \rightarrow [HClCH_3]^*$	1.93	1.78	1.79	0.68	0.92	0.10
8	$OH + C_2H_6 \rightarrow [OHC_2H_6]^*$	4.66	4.21	2.69	1.28	1.72	2.69
9	$F + H_2 \rightarrow [FH_2]^*$	3.40	3.14	1.20	0.52	0.78	1.13
10	$O + CH_4 \rightarrow [OHCH_3]^*$	3.40	3.12	2.37	0.70	1.20	13.62
11	$H + PH_3 \rightarrow [HPH_3]^*$	0.93	0.86	0.59	-0.16	0.10	2.29
12	$H + HO \rightarrow [OHH]^*$	2.03	1.59	0.44	-0.61	-0.26	10.25
13	$H + H_2S \rightarrow [HH_2S]^*$	1.01	0.92	0.65	-0.11	0.14	3.17
14	$O + HCl \rightarrow [OHCl]^*$	6.33	6.01	3.58	0.79	1.51	9.74
15	$NH_2 + CH_3 \rightarrow [CH_3NH_2]^*$	2.48	2.22	1.99	0.49	0.86	7.66
16	$NH_2 + C_2H_5 \rightarrow [NH_2C_2H_5]^*$	2.48	2.22	2.09	0.55	0.92	8.21
17	$C_2H_6 + NH_2 \rightarrow [C_2H_6NH_2]^*$	3.30	3.00	2.73	1.23	1.62	10.39
18	$NH_2 + CH_4 \rightarrow [NH_2CH_4]^*$	2.98	2.72	2.55	1.11	1.48	13.23

	Reaction Barrier	ΔCEPA_0	$\Delta\text{DC-12}$	ΔCCSD	ΔOCEPA_0	$\Delta\text{ODC-12}$	CCSD(T)
1	$[\text{HHCl}]^* \leftarrow \text{H}_2 + \text{Cl}$	1.44	1.31	1.61	0.53	0.77	7.39
2	$[\text{OHH}_2]^* \leftarrow \text{H} + \text{H}_2\text{O}$	2.09	1.66	0.09	-0.91	-0.58	21.07
3	$[\text{CH}_3\text{H}_2]^* \leftarrow \text{H} + \text{CH}_4$	0.95	0.80	0.38	-0.38	-0.11	14.91
4	$[\text{OHCH}_4]^* \leftarrow \text{CH}_3 + \text{H}_2\text{O}$	3.23	2.80	1.87	0.27	0.65	18.09
6	$[\text{OHNH}_3]^* \leftarrow \text{H}_2\text{O} + \text{NH}_2$	5.46	4.62	3.14	0.79	1.33	13.17
7	$[\text{HClCH}_3]^* \leftarrow \text{Cl} + \text{CH}_4$	1.97	1.94	2.31	0.78	1.16	5.89
8	$[\text{OHC}_2\text{H}_6]^* \leftarrow \text{H}_2\text{O} + \text{C}_2\text{H}_5$	3.34	2.89	1.85	0.28	0.64	18.49
9	$[\text{FH}_2]^* \leftarrow \text{HF} + \text{H}$	1.27	0.88	-0.78	-1.47	-1.33	32.95
10	$[\text{OHCH}_3]^* \leftarrow \text{OH} + \text{CH}_3$	2.62	2.29	1.82	0.32	0.68	7.43
11	$[\text{HPH}_3]^* \leftarrow \text{PH}_2 + \text{H}_2$	1.14	1.11	1.37	0.39	0.63	23.21
12	$[\text{OHH}]^* \leftarrow \text{H}_2 + \text{O}$	3.47	3.08	1.99	0.62	1.07	12.81
13	$[\text{HH}_2\text{S}]^* \leftarrow \text{H}_2 + \text{HS}$	1.51	1.51	1.88	0.65	0.97	16.41
14	$[\text{OHCl}]^* \leftarrow \text{OH} + \text{Cl}$	5.59	5.35	3.55	0.51	1.24	9.35
15	$[\text{CH}_3\text{NH}_2]^* \leftarrow \text{CH}_4 + \text{NH}$	2.77	2.49	2.26	0.73	1.12	21.32
16	$[\text{NH}_2\text{C}_2\text{H}_5]^* \leftarrow \text{C}_2\text{H}_6 + \text{NH}$	3.06	2.75	2.46	0.85	1.26	18.52
17	$[\text{C}_2\text{H}_6\text{NH}_2]^* \leftarrow \text{NH}_3 + \text{C}_2\text{H}_5$	2.54	2.30	2.30	0.63	1.02	16.20
18	$[\text{NH}_2\text{CH}_4]^* \leftarrow \text{CH}_3 + \text{NH}_3$	2.51	2.29	2.21	0.56	0.96	15.69
	$\Delta_{\text{MAE}}:$	2.77	2.49	1.84	0.67	0.94	
	$\Delta_{\text{SD}}:$	1.51	1.39	1.06	0.62	0.71	
	$\Delta_{\text{rel}} \text{ } \%$:	99	90	77	29	40	

largest Δ_{MAE} values come from CEPA₀ and DC-12 (2.77 and 2.49 kcal mol⁻¹, respectively). Orbital optimization greatly improves the accuracy of these methods, resulting in Δ_{MAE} values of just 0.67 and 0.94 kcal mol⁻¹ for OCEPA₀ and ODC-12, respectively. The CCSD method shows intermediate performance with $\Delta_{\text{MAE}} = 1.84$ kcal mol⁻¹. A similar trend is observed for the Δ_{SD} values, with OCEPA₀ (0.62 kcal mol⁻¹) and ODC-12 (0.71 kcal mol⁻¹) significantly improving upon CEPA₀ (1.51 kcal mol⁻¹), DC-12 (1.39 kcal mol⁻¹), and CCSD (1.06 kcal mol⁻¹). In addition to Δ_{MAE} and Δ_{SD} , table 2.2 includes mean percent error (Δ_{rel}) values, which are commonly used to benchmark performance for reaction kinetics. The smallest Δ_{rel} values are 29% and 40% for OCEPA₀ and ODC-12, respectively.

Turning to barrier heights for individual hydrogen-transfer reactions (table 2.2), the largest errors are observed for reactions 6 and 14, both involving the OH radical, for which CEPA₀ and DC-12 give errors of ~ 5 -6 kcal mol⁻¹. The best results for these reactions are obtained from OCEPA₀, with errors ranging from 0.51 to 1.18 kcal mol⁻¹. The ODC-12 method tends to predict larger barrier heights than OCEPA₀, yielding smaller errors only when OCEPA₀ underestimates the barrier heights.

2.5.3 Radical Stabilization Energies

In this section we study the performance of DCFT methods for predicting radical stabilization energies (RSEs). An R-group’s RSE is defined as the enthalpy of a

Figure 2.2: Mean absolute deviations (Δ_{MAE} , kcal mol⁻¹) and the standard deviations from the mean signed error (Δ_{SD} , kcal mol⁻¹) of barrier heights for 18 hydrogen-transfer reactions ($\text{R}_1 + \text{R}_2\text{H} \rightarrow \text{R}_1\text{H} + \text{R}_2$, HTBH38 database) computed using five methods with the aug-cc-pVTZ basis set. The errors are relative to CCSD(T)/aug-cc-pVTZ reference values. The Δ_{MAE} value is represented as a height of each colored box, while the Δ_{SD} value is depicted as a radius of the black vertical bar. See table 2.2 for data on individual database members.

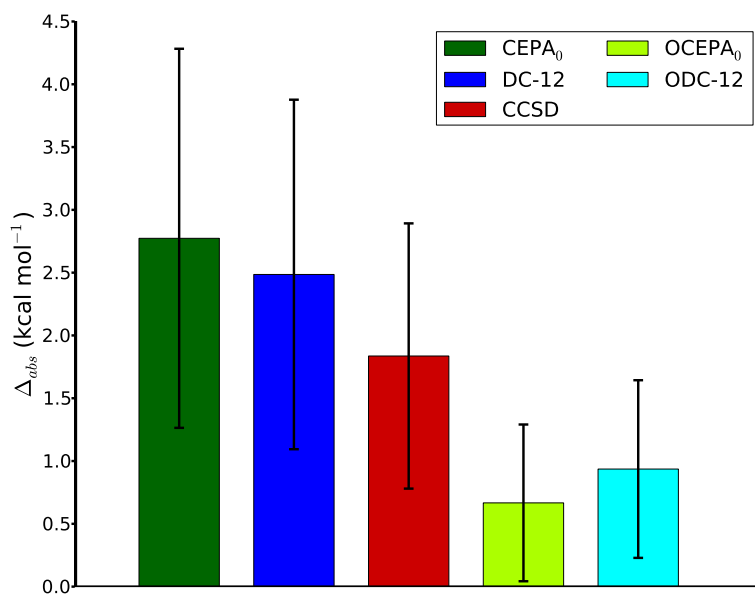
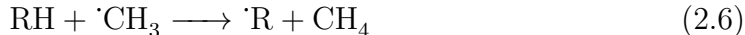


Table 2.3: Errors in radical stabilization energies (RSEs, kcal mol⁻¹) for 30 open-shell doublet species ($\cdot\text{R}$) comprising the RSE30 database⁶³ computed using six methods with the cc-pCVTZ basis set. The errors are relative to CCSD(T) with an added quadruples correction ($\delta Q = E_{\text{CCSDT(Q)}} - E_{\text{CCSD(T)}}$) shown in the rightmost column in kcal mol⁻¹. The δQ correction was computed using the cc-pCVDZ basis set. RSE is defined as the reaction enthalpy for the homodesmotic reaction $\cdot\text{CH}_3 + \text{RH} \rightarrow \text{CH}_4 + \cdot\text{R}$. To indicate the degree of spin-contamination in the UHF reference, the spin expectation values ($\langle \hat{S}^2 \rangle_{\text{SCF}}$) are also shown in units of \hbar^2 . For each method the mean absolute deviations from CCSD(T)+ δQ (Δ_{MAE} , kcal mol⁻¹) and the standard deviations from the mean signed error (Δ_{SD} , kcal mol⁻¹) are also presented.

$\cdot\text{R}$	$\langle \hat{S}^2 \rangle_{\text{SCF}}$	ΔCEPA_0	$\Delta\text{DC-12}$	ΔCCSD	ΔOCEPA_0	$\Delta\text{ODC-12}$	$\Delta\text{CCSD(T)}$	CCSD(T)+ δQ
$\cdot\text{CH}_2\text{NO}_2$	0.78	1.24	0.95	0.66	0.16	0.27	0.32	-3.50
$\cdot\text{CH}_2\text{OCHO}$	0.76	1.16	1.12	0.63	0.40	0.48	0.10	-4.84
$\cdot\text{CH}_2\text{SCH}_3$	0.76	1.89	1.70	0.81	0.63	0.72	0.15	-11.01
$\cdot\text{CF}=\text{CH}_2$	0.94	6.12	3.71	0.96	0.42	0.64	0.46	6.26
$\cdot\text{CH}_2\text{CH}_2\text{F}$	0.76	0.30	0.27	0.13	0.08	0.10	0.04	-1.53
$\cdot\text{CH}_2\text{CHO}$	0.93	5.01	2.86	0.32	-0.16	0.02	0.46	-10.11
$\cdot\text{CH}_2\text{CN}$	0.94	6.36	3.52	0.65	-0.02	0.21	0.46	-8.66
$\cdot\text{CH}_2\text{F}$	0.76	1.03	1.00	0.52	0.55	0.57	0.06	-4.22
$\cdot\text{CH}_2\text{NH}_2$	0.76	1.28	1.18	0.59	0.50	0.52	0.06	-12.06
$\cdot\text{CH}_2\text{NH}_3^+$	0.76	0.16	0.10	0.08	0.06	0.03	0.02	4.58
$\cdot\text{CH}_2\text{NHOH}$	0.77	1.76	1.57	0.78	0.58	0.64	0.15	-8.81
$\cdot\text{CH}_2\text{OH}$	0.76	1.29	1.23	0.62	0.57	0.60	0.07	-9.27
$\cdot\text{CH}_2\text{PH}_3^+$	0.76	0.21	0.14	0.01	0.01	-0.02	0.05	0.49
$\cdot\text{CH}_2\text{SH}_2^+$	0.77	0.41	0.30	0.12	0.11	0.08	0.06	2.29
$\cdot\text{CH}_2\text{SH}$	0.76	1.60	1.43	0.68	0.57	0.63	0.12	-9.68

R	$\langle \hat{S}^2 \rangle_{\text{SCF}}$	ΔCEPA_0	$\Delta\text{DC-12}$	ΔCCSD	ΔOCEPA_0	$\Delta\text{ODC-12}$	$\Delta\text{CCSD(T)}$	$\text{CCSD(T)}+\delta\text{Q}$
$\cdot\text{CH}_2\text{C}\equiv\text{CH}$	1.00	6.23	3.47	0.82	-0.03	0.23	0.52	-13.17
$\cdot\text{CH}_2\text{CH}_3$	0.76	0.30	0.26	0.11	0.08	0.10	0.03	-3.36
$\cdot\text{CH}_2\text{Cl}$	0.77	1.13	1.02	0.50	0.48	0.51	0.09	-5.67
$\cdot\text{CH}_2\text{BH}_2$	0.76	0.17	0.17	0.05	0.03	0.04	0.05	-11.66
$\cdot\text{CHO}$	0.77	2.26	2.24	1.48	1.55	1.56	0.20	-17.61
$\cdot\text{CH}_2\text{PH}_2$	0.76	1.17	1.02	0.39	0.36	0.39	0.12	-6.50
$\cdot\text{CHClF}$	0.76	1.61	1.52	0.76	0.78	0.81	0.13	-6.61
$\cdot\text{CHFCH}_3$	0.76	1.07	1.01	0.50	0.51	0.53	0.08	-5.87
$\cdot\text{CH(OH)}_2$	0.76	1.30	1.22	0.60	0.60	0.61	0.08	-6.67
$\cdot\text{CHCl}_2$	0.77	1.78	1.57	0.72	0.72	0.75	0.15	-9.56
$\cdot\text{CHF}_2$	0.76	1.50	1.48	0.78	0.83	0.85	0.10	-4.07
$\text{CH}_2=\text{C}\cdot-\text{CN}$	1.39	19.10	11.50	2.36	-0.31	0.29	1.80	1.98
$\cdot\text{C}\equiv\text{CH}$	1.15	11.20	6.51	0.77	-0.78	-0.07	0.82	26.25
$\cdot\text{CH}=\text{CH}_2$	0.94	5.42	3.01	0.58	0.11	0.31	0.40	5.49
$\cdot\text{CH}_2-\text{CH}=\text{CH}_2$	0.97	4.98	3.17	0.51	0.11	0.31	0.48	-17.53
		$\Delta_{\text{MAE}}:$	2.97	2.01	0.62	0.40	0.43	0.25
		$\Delta_{\text{SD}}:$	3.97	2.27	0.45	0.43	0.35	0.35

homodesmotic reaction



where exothermic (negative) values indicate that the radical $\cdot\text{R}$ is more thermodynamically stable than $\cdot\text{CH}_3$.⁶⁵ For our benchmark we use the RSE30 dataset⁶³, which provides a diverse variety of $\cdot\text{R}$ species (listed in table 2.3). Since the performance of CCSD(T) is known to deteriorate for strongly spin-contaminated UHF references,^{66–70} we augment CCSD(T) energies with a quadruples correction ($\Delta\text{Q} = E_{\text{CCSDT(Q)}} - E_{\text{CCSD(T)}}$) and use these as our benchmark. CBS-extrapolated CCSD(T) reference values have been published for this dataset,³¹ but we use CCSD(T) values computed with the cc-pCVTZ basis set to avoid basis-set incompleteness effects. The δQ correction was evaluated using the cc-pCVDZ basis set. As in the previous Section, DC-06 and ODC-06 computations cannot be converged for all database members and are omitted in the analysis below (see Supporting Information for incomplete DC-06 and ODC-06 data).

The relative performance of the DCFT, CEPA, and CC methods for the RSE30 dataset is shown in fig. 2.3. The effect of orbital-optimization on accuracy is now even more pronounced, reducing the large Δ_{MAE} errors of CEPA_0 (2.97 kcal mol⁻¹) and DC-12 (2.01 kcal mol⁻¹) to 0.40 and 0.43 kcal mol⁻¹ for OCEPA_0 and ODC-12, respectively. CCSD has a slightly larger Δ_{MAE} value (0.62 kcal mol⁻¹), while CCSD(T) has the smallest overall Δ_{MAE} (0.25 kcal mol⁻¹). Both CEPA_0 and DC-12 show large standard deviations again (3.97 and 2.27 kcal mol⁻¹, respectively).

For OCEPA₀, the standard deviation (0.43 kcal mol⁻¹) is similar to that of CCSD (0.45 kcal mol⁻¹). ODC-12 and CCSD(T) exhibit the most consistent performance with the same Δ_{SD} value of 0.35 kcal mol⁻¹.

Deviations from CCSD(T)+ δ Q for individual RSEs predicted by each method are tabulated in table 2.3. In addition, table 2.3 includes expectation values of the square-norm spin operator computed for the UHF wavefunction of R ($\langle \hat{S}^2 \rangle_{\text{SCF}}$). The largest errors in computed RSEs were obtained for R species with $\langle \hat{S}^2 \rangle_{\text{SCF}} > 0.9 \hbar^2$ (radicals 4, 6, 7, 16, and 27-30 in table 2.3). For these systems, the average CEPA₀ and DC-12 errors are 8.05 and 4.72 kcal mol⁻¹, and the average CCSD(T) error is 0.68 kcal mol⁻¹. OCEPA₀ and ODC-12 offer remarkably better performance for this subset, with average errors of 0.24 kcal mol⁻¹ and 0.26 kcal mol⁻¹.

2.5.4 Adiabatic Ionization Energies in Electron-Dense Molecules

We conclude the assessment of DCFT methods for the description of thermodynamic properties by computing adiabatic ionization energies (AIEs) for a set of 10 di- and triatomic electron-dense molecules (table 2.4), i.e. those that are composed of elements with small atomic radius, high electron affinity, and high electronegativity (N, O, F), in order to increase the magnitude of electron correlation effects. We use experimentally measured ionization energies reported to high precision (~ 0.01 eV)⁷¹⁻⁷⁴ as reference values for our benchmark (IE_{ref}, table 2.4). The AIEs were computed using the cc-pCVQZ basis set, with harmonic ZPVE corrections applied to each neutral and cationic system.

Figure 2.3: Mean absolute deviations (Δ_{MAE} , kcal mol⁻¹) and the standard deviations from the mean signed error (Δ_{SD} , kcal mol⁻¹) of the radical stabilization energies (RSEs) for 30 open-shell doublet species (RSE30 database) computed using six methods with the cc-pCVTZ basis set. The errors are relative to CCSD(T) with an added quadruples correction ($\delta Q = E_{\text{CCSDT(Q)}} - E_{\text{CCSD(T)}}$). The δQ correction was computed using the cc-pCVDZ basis set. RSE is defined as the reaction enthalpy for the homodesmotic reaction $\cdot\text{CH}_3 + \text{RH} \rightarrow \text{CH}_4 + \cdot\text{R}$. The Δ_{MAE} value is represented as a height of each colored box, while the Δ_{SD} value is depicted as a radius of the black vertical bar. See table 2.3 for data on individual database members.

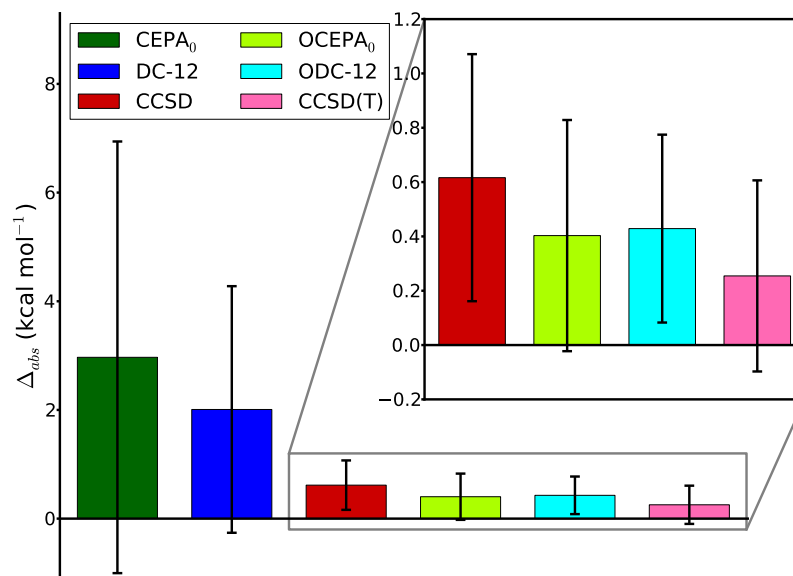


Table 2.4: Errors in adiabatic ionization energies (AIEs, eV) for 10 di- and triatomic molecules computed using six methods with the cc-pCVQZ basis set. The errors are relative to experimental values (IE_{ref} , eV) from Ref. 71, unless noted otherwise. For all AIEs the harmonic zero-point vibrational energy corrections were included. For each method the mean absolute deviations from IE_{ref} (Δ_{MAE} , eV) and the standard deviations from the mean signed error (Δ_{SD} , eV) are also shown.

Molecule	Transition	ΔCEPA_0	$\Delta\text{DC-12}$	ΔCCSD	ΔOCEPA_0	$\Delta\text{ODC-12}$	$\Delta\text{CCSD(T)}$	IE_{ref}
N ₂	$^1\Sigma_g^+ \rightarrow ^2\Sigma_g^+$	0.08	0.17	0.12	-0.05	0.07	-0.03	15.581 ± 0.008^a
O ₂	$^3\Sigma_g^- \rightarrow ^2\Pi_g$	-0.11	-0.03	0.04	-0.09	-0.02	-0.04	12.0697 ± 0.0002
F ₂	$^1\Sigma_g^+ \rightarrow ^2\Pi_g$	0.06	0.06	0.04	0.08	0.01	-0.03	15.697 ± 0.003
NO	$^2\Pi \rightarrow ^1\Sigma^+$	-0.15	-0.05	-0.05	-0.05	-0.02	-0.09	9.26438 ± 0.00005
OF	$^2\Pi \rightarrow ^3\Sigma^-$	0.11	0.12	-0.10	-0.03	-0.02	-0.11	12.77 ± 0.01^b
HNC	$^1\Sigma_g^+ \rightarrow ^2\Sigma^+$	0.27	0.14	-0.12	-0.14	-0.08	-0.04	12.04 ± 0.01^c
HOF	$^1A' \rightarrow ^2A''$	0.20	0.17	-0.10	-0.03	-0.04	-0.07	12.71 ± 0.01
FNO	$^1A' \rightarrow ^2A''$	0.51	0.10	-0.02	-0.02	-0.00	0.04	12.63 ± 0.03
F ₂ N	$^2B_1 \rightarrow ^1A_1$	0.07	0.10	0.07	0.01	0.03	-0.08	11.63 ± 0.01
F ₂ O	$^1A_1 \rightarrow ^2B_1$	0.49	0.37	-0.01	0.05	0.04	-0.04	13.11 ± 0.01
$\Delta_{\text{MAE}}:$		0.21	0.13	0.06	0.05	0.03	0.06	
$\Delta_{\text{SD}}:$		0.22	0.12	0.08	0.06	0.04	0.04	

^a Reference 72.

^b Reference 73.

^c Reference 74.

The Δ_{MAE} and Δ_{SD} values for our computed AIEs relative to experiment are plotted in fig. 2.4. Of the six methods, CEPA₀ and DC-12 exhibit the largest Δ_{MAE} values (0.21 and 0.13 eV, respectively). The closest agreement with experiment is given by ODC-12, with $\Delta_{\text{MAE}} = 0.03$ eV. OCEPA₀, CCSD, and CCSD(T) show somewhat poorer performance ($\Delta_{\text{MAE}} = 0.05, 0.06$ and 0.06 eV, respectively). The Δ_{SD} for ODC-12 matches that of CCSD(T) (0.04 eV). For the other methods, the Δ_{SD} values decrease in the order CEPA₀ (0.22 eV) > DC-12 (0.12) > CCSD (0.08) > OCEPA₀ (0.06).

Individual errors for each system are shown in table 2.4. Both DC-12 and CEPA₀ exhibit large deviations for F₂O (0.49 and 0.37 eV), and CEPA₀ also gives a large error for FNO (0.51 eV) which is the maximum error for this dataset. Both DC-12 and CEPA₀ give errors exceeding 0.1 eV for seven of the ten systems, whereas CCSD exhibits errors in excess of 0.1 eV for only three systems (OF, HNC, and HOF). CCSD(T) has only one such error (0.11 eV for OF), as does OCEPA₀ (0.14 eV for HNC). ODC-12 does the best of the methods considered, with a maximum error of 0.08 eV, found for the AIE of HNC.

2.5.5 Covalent Bond Stretching in Diatomic Molecules

Finally, we benchmark DCFT methods for covalent bond stretching. Although accurate description of bond stretching demands the use of multireference methods, our aim here is to explore the limits of DCFT away from equilibrium. For this purpose, we compute the energy as a function of bond distance for diatomic molecules with single (HF and BH), double (BeO), and triple (N₂) bonds using the

Figure 2.4: Mean absolute deviations (Δ_{MAE} , eV) and the standard deviations from the mean signed error (Δ_{SD} , eV) of adiabatic ionization energies for 10 di- and triatomic molecules computed using six methods with the cc-pCVQZ basis set. The errors are relative to experimental values.^{71–74} The Δ_{MAE} value is represented as a height of each colored box, while the Δ_{SD} value is depicted as a radius of the black vertical bar. See table 2.4 for data on individual molecules.

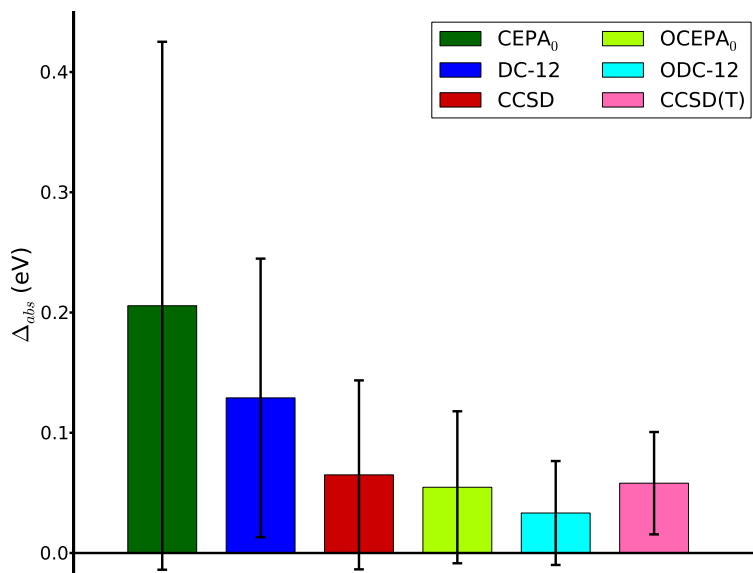


Figure 2.5: Error in the total energy (mE_h), relative to full CI, as a function of B–H internuclear separation (\AA) computed using six methods with the DZP basis set. The full CI reference is depicted with a horizontal dotted line. The dashed vertical line indicates the full CI equilibrium bond distance.

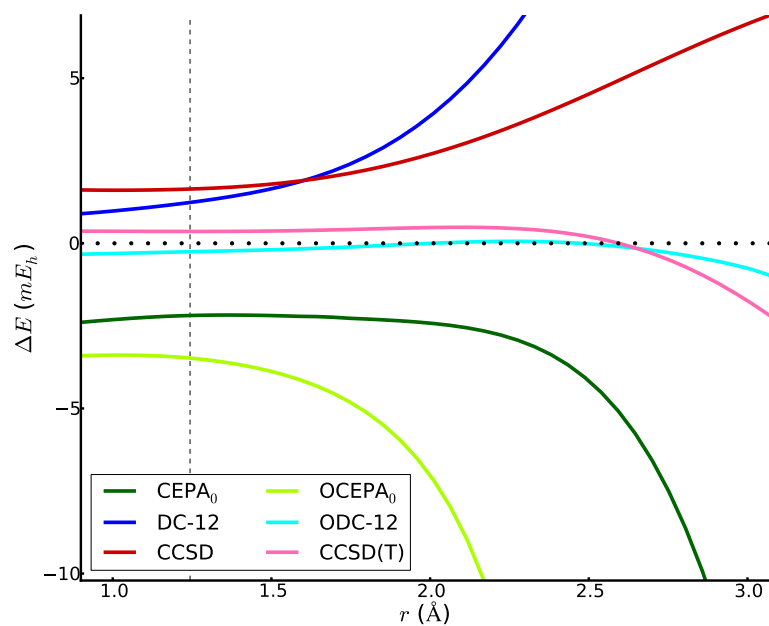


Figure 2.6: Error in the total energy (mE_h), relative to full CI, as a function of H–F internuclear separation (\AA) computed using six methods with the DZP basis set. The full CI reference is depicted with a horizontal dotted line. The dashed vertical line indicates the full CI equilibrium bond distance.

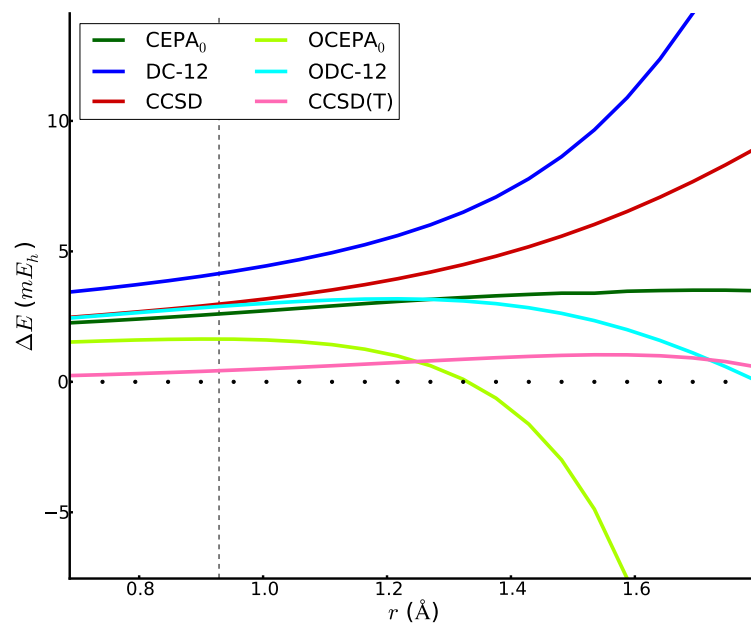


Figure 2.7: Error in the total energy (mE_h), relative to full CI, as a function of Be–O internuclear separation (\AA) computed using six methods with the 6-31G basis set. The full CI reference is depicted with a horizontal dotted line. The dashed vertical line indicates the full CI equilibrium bond distance.

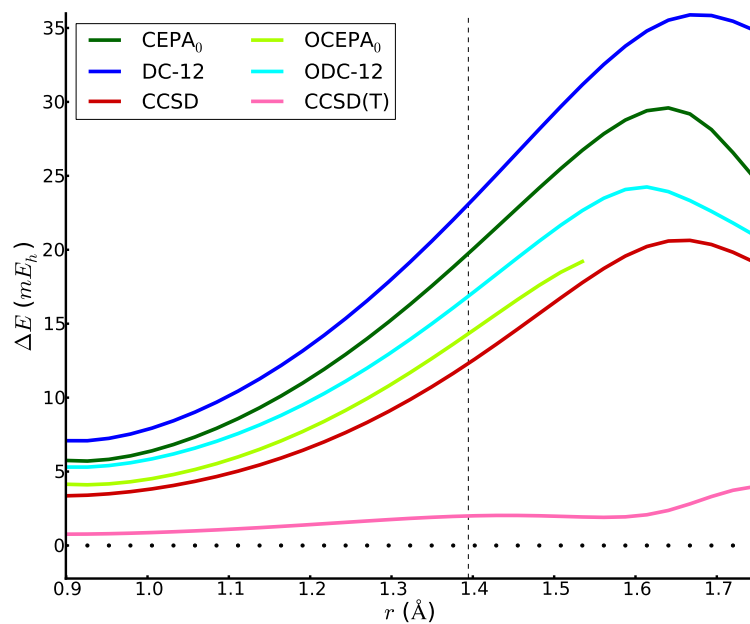
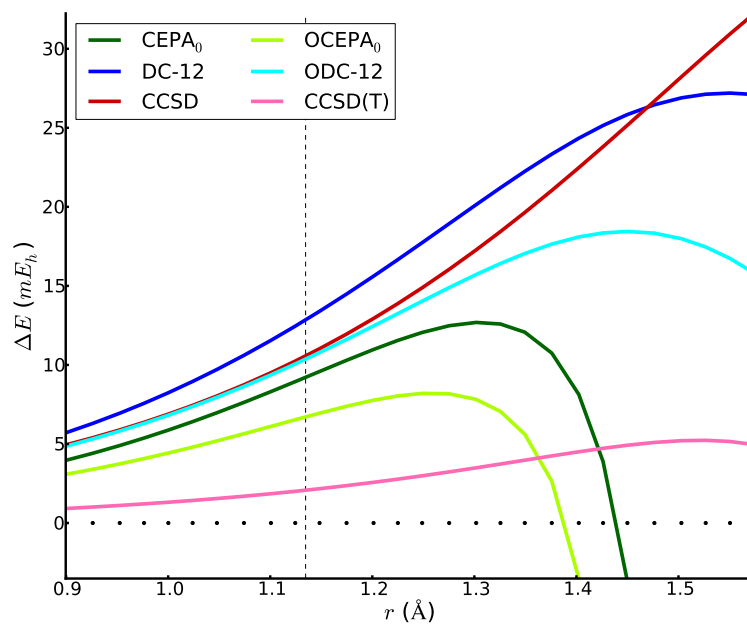


Figure 2.8: Error in the total energy (mE_h), relative to full CI, as a function of N–N internuclear separation (\AA) computed using six methods with the 6-31G basis set. The full CI reference is depicted with a horizontal dotted line. The dashed vertical line indicates the full CI equilibrium bond distance.



CEPA₀, OCEPA₀, DC-12, ODC-12, CCSD, and CCSD(T) methods. We restrict ourselves to modest basis sets in order to use full CI (FCI) as a reference, and plot the errors with respect to FCI (ΔE) as a function of internuclear distance for each molecule. The relative performance of the methods is described below using non-parallelity errors ($\text{NPE} = \Delta E_{\text{max}} - \Delta E_{\text{min}}, \text{m}E_{\text{h}}$) computed for specific bond distance ranges.

BH Figure 2.5 shows errors relative to FCI for the BH molecule. DC-12 and CCSD increasingly overestimate the energy at larger internuclear distances, whereas the CEPA₀ error curve is concave down. Orbital optimization lowers the binding energy for OCEPA₀ even further compared to CEPA₀, leading to large errors with respect to FCI for $r(\text{B-H}) \geq 1.5 r_{\text{e}}$, where r_{e} is the FCI equilibrium bond distance ($r_{\text{e}} = 1.244 \text{ \AA}$). At $1.87 r_{\text{e}}$, OCEPA₀ encounters convergence problems, which originate from numerical instabilities due to the method’s deficiencies in the description of N -representability. The ODC-12 method exhibits much more stable behavior with respect to bond stretching in this case, fortuitously showing smaller errors and better parallelity than CCSD(T). For the range $[0.72 r_{\text{e}}, 2.47 r_{\text{e}}]$, the NPEs decrease in the order DC-12 ($24 \text{ m}E_{\text{h}}$) > CEPA₀ (15) > CCSD (5) > CCSD(T) (3) > ODC-12 (1).

HF Errors for HF bond stretching are plotted in fig. 2.6. The ΔE values of CCSD and DC-12 increase as a function of $r(\text{H-F})$, while CEPA₀ fortuitously maintains parallelity similar to CCSD(T) over the range $[0.74 r_{\text{e}}, 1.94 r_{\text{e}}]$ ($r_{\text{e}} = 0.929 \text{ \AA}$). OCEPA₀ increasingly overestimates the HF binding energy away from equilibrium,

failing to converge past $1.82 r_e$. The ODC-12 method exhibits larger NPE than was observed for BH, and encounters convergence problems past $1.94 r_e$. CCSD(T) shows the best overall performance, with errors between 0 and $1 mE_h$. In the range $[0.74 r_e, 1.94 r_e]$ the computed NPE values are: DC-12 ($15 mE_h$) > CCSD (7) > ODC-12 (3) > CEPA₀ (1) \approx CCSD(T) (1). Recently, the orbital-optimized variants of CCSD(T) have been shown to yield good performance for HF bond stretching.⁷⁵

BeO The double bond of BeO presents a more challenging test for the single-reference methods under consideration (fig. 2.7). All methods but CCSD(T) show qualitatively similar error curves, with inflection points near the FCI equilibrium ($r_e = 1.394 \text{ \AA}$) and valleys/peaks around $0.6 r_e/1.2 r_e$. OCEPA₀ encounters convergence problems past $1.10 r_e$. The ODC-12 method performs similarly to CCSD. Overall, the NPEs for the range $[0.65 r_e, 1.10 r_e]$ decrease in the following order: DC-12 ($29 mE_h$) > CEPA₀ (24) > ODC-12 (19) > CCSD (17) > CCSD(T) (3).

N₂ Figure 2.8 depicts the errors relative to FCI for triple bond stretching in N₂. Here, OCEPA₀ fails to converge past $1.24 r_e$ ($r_e = 1.135 \text{ \AA}$). The ODC-12 method significantly overestimates the binding energy, possibly due to the lack of three-body correlation effects, but shows much more stable performance compared to methods other than CCSD(T). NPEs in the range $[0.79 r_e, 1.39 r_e]$ decrease in the order: CEPA₀ ($802 mE_h$)[‡] > CCSD (27) > DC-12 (21) > ODC-12 (14) > CCSD(T) (4).

[‡]CEPA₀ exhibits a vertical asymptote at $1.36 r_e$ for N₂ stretching.

2.6 Conclusions

We have presented the benchmark study of four density cumulant functional theory (DCFT) methods (DC-06, DC-12, ODC-06, and ODC-12) developed recently in our group.^{49;50;52;53} Specifically we have compared the performance of DCFT to that of coupled electron pair methods (CEPA₀ and OCEPA₀), as well as coupled-cluster theory [CCSD and CCSD(T)] for predicting a variety of chemical properties relevant to thermochemistry and kinetics, with a particular focus on open-shell, electron-dense, and non-equilibrium systems.

Our results indicate that among the four DCFT methods, the best agreement with available reference data is obtained for the ODC-12 method. While all four DCFT formulations yield similar results for the description of noncovalent interactions, DC-06, DC-12, and ODC-06 exhibit worse performance than ODC-12 for thermodynamic and kinetic properties of reactions involving open-shell molecules. In particular, DC-06 and ODC-06 frequently encounter convergence problems that originate from poor description of N -representability. In comparing ODC-12 to other methods, several trends can be observed:

(i) For all benchmark datasets, ODC-12 outperforms CCSD with errors smaller by almost a factor of two, on average. ODC-12 is also superior to CCSD for the description of single bond stretching in BH and HF, although it does not converge for all bond distances.

(ii) The performance of ODC-12 and OCEPA₀ is comparable. In particular, for hydrogen-transfer reaction barrier heights, the OCEPA₀ method yields smaller percent errors than ODC-12, whereas, for the radical stabilization energies (RSE)

and adiabatic ionization energies (AIE) in electron-dense molecules, the ODC-12 method smaller standard deviations than OCEPA₀. For AIEs, ODC-12 gives smaller mean absolute deviations by almost a factor of two. ODC-12 also shows significantly smaller non-parallelity errors than OCEPA₀ for covalent bond stretching, and can be converged for a larger range of distances for all diatomic molecules studied.

(iii) For the two most challenging datasets, RSE and AIE, the standard deviation of ODC-12 and CCSD(T) are similar. While CCSD(T) yields smaller mean absolute errors for the RSE database, the ODC-12 method significantly outperforms CCSD(T) for the AIE test case. However, for bond stretching ODC-12 is competitive with CCSD(T) only for the BH dissociation and shows worse results for other molecules.

Overall, the data presented herein indicates that the ODC-12 method can be used as an efficient $\mathcal{O}(n^6)$ alternative to CCSD, capable of predicting thermodynamic and kinetic quantities that are competitive in accuracy with the “gold-standard” $\mathcal{O}(n^7)$ CCSD(T). Although our current implementation of ODC-12 is far from optimal, the ODC-12 equations have reduced non-linearities compared to CCSD, which makes them more amenable to parallel implementation. The efficiency of ODC-12 can also greatly benefit from spin-adaptation,^{42;76;77} local approximations,^{34;78–80} and density fitting.^{78;81–83} Another important advantage of ODC-12 over CCSD is its stationarity, which makes the computation of first-order properties and analytic gradients more efficient and easily accessible. In particular, ODC-12 has potential to be used for computing accurate response properties

which do not suffer from a lack of gauge-invariance.^{1;84}

Chapter 3

Linear-Response Density Cumulant Theory for Excited States: First Implementation and Benchmark Calculations*

3.1 Abstract

We present a linear-response formulation of density cumulant functional theory (DCT) that provides accurate access to many electronic states. DCT expresses the electronic energy as a Hermitian, size-extensive, and stationary functional of the one-particle density matrix and the two-particle density cumulant. In the original DCT formulation only the information about a single electronic state (usually, the ground state) is obtained. In this research, we combine DCT with linear response theory to obtain information about many electronic states simultaneously. We discuss the derivation of linear-response DCT, present its implementation for the ODC-12 method (LR-ODC-12), and benchmark its performance against highly accurate equation-of-motion coupled cluster theory with up to full triple excitations (EOM-CCSDT). Our results for a set of small molecules demonstrate that LR-ODC-12 vertical excitation energies are in closer agreement with EOM-CCSDT

*A. V. Copan and A. Yu. Sokolov (to be submitted in J. Chem. Theory Comput).

than those obtained from equation-of-motion coupled cluster theory with up to double excitations (EOM-CCSD). In addition, we report a linear-response formulation of the orbital-optimized linearized coupled cluster theory with double excitations (LR-OLCCD), which we obtain by neglecting the non-linear terms in the LR-ODC-12 equations.

3.2 Introduction

Accurate simulation of excited electronic states remains one of the major challenges in modern electronic structure theory. *Ab initio* methods for excited states can be divided into single-reference and multi-reference categories, based on their ability to treat static electron correlation. Multi-reference methods can correctly describe static correlation in near-degenerate valence orbitals and electronic states with multiple-excitation character, but often lack accurate treatment of important dynamic correlation effects (e.g., multi-configurational self-consistent field or multi-reference perturbation theories)^{85–94} or become very costly when the number of near-degenerate orbitals is large (e.g., multi-reference configuration interaction or coupled cluster theories).^{95–107} Meanwhile, single-reference methods^{21;23;108–120} often provide a compromise between the computational cost and accuracy, and can be used to reliably compute properties of molecules in low-lying electronic states near the equilibrium geometries. In these situations, single-reference equation-of-motion coupled cluster theory (EOM-CC)^{21;23;110–113} is usually the method of choice, especially when high accuracy is desired.

The EOM-CC methods yield size-intensive excitation energies^{115;116} and can be

systematically improved by increasing the excitation rank of the cluster operator in the exponential parametrization of the wavefunction. Although EOM-CC is usually formulated in the context of a similarity-transformed Hamiltonian, its excitation energies are equivalent to those obtained from linear-response coupled cluster theory (LR-CC).^{114–116} Both EOM-CC and LR-CC are based on non-Hermitian eigenvalue problems, complicating the computation of molecular properties (e.g., transition dipoles) by requiring evaluation of left and right eigenvectors.^{121–124} Several Hermitian alternatives to EOM-CC and LR-CC have been proposed to avoid these problems, such as algebraic diagrammatic construction^{125–129}, unitary and variational LR-CC,^{130–132} similarity-constrained CC,¹³³ and propagator-based LR-CC.^{134;135}

In this work, we present the development of linear-response density cumulant functional theory (LR-DCT), a size-intensive approach for excited electronic states. In density cumulant functional theory (DCT),^{35;49;50;52;53;136;137} the electronic energy is obtained by optimizing the energy functional directly in terms of the one-particle reduced density matrix and the two-particle density cumulant, a fully connected part of the two-particle reduced density matrix (2-RDM).^{39–42;44;48;138–141} In this regard, DCT is related to approaches that are based on the variational optimization^{41;142–147} or parametrization^{36–38} of 2-RDM. On the other hand, DCT has a close relationship^{52;53} with wavefunction-based electronic structure theories, such as linearized, unitary, and variational coupled cluster theory.^{5;148–155} In contrast to variational 2-RDM theory^{45–47} and traditional coupled cluster methods [e.g., CCSD and CCSD(T)],^{21;23} DCT naturally combines size-extensivity and a

Hermitian energy functional. In addition, the DCT electronic energy is fully relaxed with respect to all of its parameters, which greatly simplifies computation of the first-order molecular properties.^{156–159} We have successfully applied DCT to a variety of chemical systems with different electronic structure effects (e.g., open-shell, symmetry-breaking, and multi-reference).^{53;136;137;160;161} One limitation of the original DCT formulation is that it can only obtain information about the lowest-energy state of a particular symmetry (usually, the ground state). By combining DCT with linear response theory, we remove this limitation, providing access to many electronic states simultaneously.

We begin with a brief overview of DCT (section 3.3.1) and linear response theory (section 3.3.2). We then discuss the derivation of linear-response theory for the ODC-12 method (LR-ODC-12, section 3.3.3). In section 3.3.4, we derive equations for the linear-response orbital-optimized linearized coupled cluster theory with double excitations (LR-OLCCD), which we obtain by neglecting the non-linear terms in the LR-ODC-12 equations. We outline the computational details in section 3.4. In section ??, we demonstrate that the LR-ODC-12 excitation energies are size-intensive (??), test the performance of LR-ODC-12 for the dissociation of H_2 (??), and benchmark the accuracy of LR-ODC-12 for vertical excitation energies of small molecules (??). Finally, we present our conclusions in section ??.

3.3 Theory

3.3.1 Overview of Density Cumulant Theory

We begin with a brief overview of DCT for a single electronic state. Our starting point is to express the electronic energy as a trace of the one- and antisymmetrized two-electron integrals (h_p^q and \bar{g}_{pq}^{rs}) with the reduced one- and two-body density matrices (γ_q^p and γ_{rs}^{pq}):

$$E = h_p^q \gamma_q^p + \frac{1}{4} \bar{g}_{pq}^{rs} \gamma_{rs}^{pq} \quad (3.1)$$

where summation over the repeated indices is implied. In DCT, the two-body density matrix γ_{rs}^{pq} is expanded in terms of its connected part, the two-body density cumulant (λ_{rs}^{pq}), and its disconnected part, which is given by an antisymmetrized product of one-body density matrices:³⁵

$$\gamma_{rs}^{pq} = \langle \Psi | a_{rs}^{pq} | \Psi \rangle = \lambda_{rs}^{pq} + P_{(r/s)} \gamma_r^p \gamma_s^q \quad (3.2)$$

where $P_{(r/s)} v_{rs} = v_{rs} - v_{sr}$ denotes antisymmetrization and $a_{rs}^{pq} = a_p^\dagger a_q^\dagger a_s a_r$ is the two-body operator in second quantization. The one-body density matrix γ_q^p is determined from its non-linear relationship to the cumulant's partial trace:⁵²

$$\gamma_q^p = \gamma_r^p \gamma_q^r - \lambda_{qr}^{pr} \quad (3.3)$$

This reduces eq. (3.1) to a functional of the two-body cumulant and the basis of spin-orbitals, thereby defining the DCT energy functional. The density cumulant

is parametrized by choosing a specific Ansatz for the wavefunction $|\Psi\rangle$ such that¹³⁶

$$\lambda_{rs}^{pq} = \langle \Psi | a_{rs}^{pq} | \Psi \rangle_c \quad (3.4)$$

where c indicates that only fully connected terms are included in the parametrization. eq. (3.4) can be considered as a set of n -representability conditions that ensure that the resulting one- and two-electron density matrices represent a physical n -electron wavefunction. To compute the DCT energy, the functional (3.1) is made stationary with respect to all of its parameters. Importantly, due to the connected nature of eq. (3.4), DCT is both size-consistent and size-extensive for any parametrization of $|\Psi\rangle$, and is exact in the limit of a complete parametrization.¹³⁶

In this work, we consider the ODC-12 method,^{52;53} which parametrizes the cumulant through a unitary treatment of single excitations and a linear expansion of double excitations.

$$|\Psi\rangle = e^{T_1 - T_1^\dagger} (1 + \hat{T}_2) |\Phi\rangle \quad (3.5)$$

$$T_1 = \mathbf{t}_1 \cdot \mathbf{a}_1 = t_a^i a_i^a \quad (3.6)$$

$$T_2 = \mathbf{t}_2 \cdot \mathbf{a}_2 = \frac{1}{4} t_{ab}^{ij} a_{ij}^{ab} \quad (3.7)$$

(missing hats for the operators in many places. I generally prefer to keep hats in the equations) The exponential singles operator $e^{T_1 - T_1^\dagger}$ has the effect of a unitary transformation of the spin-orbital basis and is incorporated in our implementation of the ODC-12 method by optimizing the orbitals.⁵³ The \mathbf{t}_1 and \mathbf{t}_2 parameters are

obtained from the stationarity conditions

$$\frac{\partial E}{\partial \mathbf{t}_1^\dagger} \stackrel{!}{=} 0, \quad \frac{\partial E}{\partial \mathbf{t}_2^\dagger} \stackrel{!}{=} 0 \quad (3.8)$$

and are used to compute the ODC-12 energy. Explicit equations for the stationarity conditions are given in Refs. 52 and 53. Although in ODC-12 the wavefunction parametrization is linear with respect to double excitations (eq. (3.5)), the ODC-12 energy stationarity conditions are non-linear in \mathbf{t}_2 due to the non-linear relationship between the one-particle density matrix and the density cumulant (eq. (3.3)).⁵² Neglecting the non-linear \mathbf{t}_2 terms in eq. (3.8) results in the equations that define the linearized orbital-optimized coupled cluster doubles method (OLCCD). This method is equivalent to the orbital-optimized coupled electron pair approximation zero (OCEPA₀).³⁰

3.3.2 Linear Response Theory

We now briefly review linear response theory in the quasi-energy formulation.¹⁶² The quasi-energy of a system perturbed by a time-dependent interaction $\hat{V}f(t)$ is defined as

$$Q(t) = \langle \Psi(t) | \hat{H} + \hat{V}f(t) - i \frac{\partial}{\partial t} | \Psi(t) \rangle \quad (3.9)$$

where $\Psi(t)$ is the phase-isolated wavefunction, from which the usual Schrödinger wavefunction can be recovered as $e^{-i \int_0^t dt' Q(t')} \Psi(t)$. Assuming that the perturbation is periodic

$$f(t) = \sum_{\omega} f(\omega) e^{-i\omega t} \quad (3.10)$$

the time average of the quasi-energy over a period of oscillation, denoted as $\{Q(t)\}$, is variational with respect to the exact dynamic state.¹⁶³ The independent parameters $\mathbf{u}(t)$ that define such a state can be written using a Fourier expansion

$$\mathbf{u}(t) = \sum_{n=0}^{\infty} \sum_{\omega_1 \dots \omega_n} \mathbf{u}(\omega_1, \dots, \omega_n) e^{-i(\omega_1 + \dots + \omega_n)t} \quad (3.11)$$

where the outer sum runs over polynomial orders in $f(t)$. The stationarity of the time-averaged quasi-energy then implies the following relationship¹⁶⁴

$$0 = \left. \frac{d}{df(\omega)} \frac{\partial \{Q(t)\}}{\partial \mathbf{u}^\dagger(\omega)} \right|_{f=0} = \left. \frac{\partial^2 \{Q(t)\}}{\partial \mathbf{u}^\dagger(\omega) \partial \mathbf{u}(\omega)} \frac{\partial \mathbf{u}(\omega)}{\partial f(\omega)} \right|_{f=0} + \left. \frac{\partial^2 \{Q(t)\}}{\partial \mathbf{u}^\dagger(\omega) \partial f(\omega)} \right|_{f=0} \quad (3.12)$$

which constitutes a linear equation for the first-order response of the system to the perturbation. When the frequency ω is in resonance with an excitation energy of the system, eq. (3.12) will result in an infinite first-order response $\frac{\partial \mathbf{u}(\omega)}{\partial f(\omega)}$. From eq. (3.12), we find that these poles occur when the Hessian matrix of the quasi-energy with respect to the wavefunction parameters $\mathbf{u}(\omega)$ becomes singular. We can express this Hessian matrix in the form

$$\left. \frac{\partial^2 \{Q(t)\}}{\partial \mathbf{u}^\dagger(\omega) \partial \mathbf{u}(\omega)} \right|_{f=0} \equiv \mathbf{E} - \omega \mathbf{M} \quad (3.13)$$

where \mathbf{E} is the Hessian of the time-averaged electronic energy $\{\langle \Psi(t) | \hat{H} | \Psi(t) \rangle\}$ and $\omega \mathbf{M}$ is the Hessian of the time-derivative overlap $\{\langle \Psi(t) | i\dot{\Psi}(t) \rangle\}$. The excitation

energies of the system ω_k can therefore be determined by solving the following generalized eigenvalue equation:

$$\mathbf{E}\mathbf{z}_k = \omega_k \mathbf{M}\mathbf{z}_k \quad (3.14)$$

where \mathbf{M} serves as the metric matrix. eq. (3.14) allows the determination of excitation energies for an arbitrary parametrization of $|\Psi(t)\rangle$.

The generalized eigenvectors \mathbf{z}_k can be used to compute transition properties for excited states. In particular, in the exact linear response theory¹⁶⁵ the transition strength of the perturbing interaction, $|\langle\Psi|\hat{V}|\Psi_k\rangle|^2$, is equal to the complex residue of the following quantity at $\omega \rightarrow \omega_k$:

$$\langle\langle\hat{V};\hat{V}\rangle\rangle_\omega \equiv \mathbf{v}'^\dagger \cdot \frac{\partial \mathbf{u}(\omega)}{\partial f(\omega)} \Big|_{f=0} \quad (3.15)$$

This quantity is known as the linear response function and \mathbf{v}' is termed the property gradient vector,¹⁶⁶ which is defined as follows:

$$\mathbf{v}' \equiv \frac{\partial^2 \{Q(t)\}}{\partial \mathbf{u}^\dagger(\omega) \partial f(\omega)} \Big|_{f=0} \quad (3.16)$$

Substituting eqs. (3.13) and (3.16) into eq. (3.12) and decomposing the quasi-energy Hessian as

$$\mathbf{E} - \omega \mathbf{M} = (\mathbf{Z}^\dagger)^{-1} (\mathbf{Z}^\dagger \mathbf{M} \mathbf{Z}) (\mathbf{\Omega} - \omega \mathbf{1}) (\mathbf{Z})^{-1} \quad (3.17)$$

where $\mathbf{\Omega}$ is a diagonal matrix of generalized eigenvalues and \mathbf{Z} is a matrix of

generalized eigenvectors that simultaneously diagonalizes \mathbf{E} and \mathbf{M} , we obtain the general formula for the transition strengths:

$$\lim_{\omega \rightarrow \omega_k} (\omega - \omega_k) \langle\langle \hat{V}; \hat{V} \rangle\rangle_{\omega} = \frac{|\mathbf{z}_k^{\dagger} \mathbf{v}'|^2}{\mathbf{z}_k^{\dagger} \mathbf{M} \mathbf{z}_k} \quad (3.18)$$

In section 3.3.3, we will use the quasi-energy formalism to derive equations for the linear-response ODC-12 method (LR-ODC-12).

3.3.3 Linear-Response ODC-12

In the ODC-12 method, the electronic energy Hessian can be written in the following form

$$\mathbf{E} = \begin{pmatrix} \mathbf{A}_{11} & \mathbf{A}_{12} & \mathbf{B}_{11} & \mathbf{B}_{12} \\ \mathbf{A}_{21} & \mathbf{A}_{22} & \mathbf{B}_{21} & \mathbf{B}_{22} \\ \mathbf{B}_{11}^* & \mathbf{B}_{12}^* & \mathbf{A}_{11}^* & \mathbf{A}_{12}^* \\ \mathbf{B}_{21}^* & \mathbf{B}_{22}^* & \mathbf{A}_{21}^* & \mathbf{A}_{22}^* \end{pmatrix} \quad (3.19)$$

where the submatrices are defined in general as

$$\mathbf{A}_{nm} = \left. \frac{\partial^2 E}{\partial \mathbf{t}_n^{\dagger} \partial \mathbf{t}_m} \right|_{f=0}, \quad \mathbf{B}_{nm} = \left. \frac{\partial^2 E}{\partial \mathbf{t}_n^{\dagger} \partial \mathbf{t}_m^*} \right|_{f=0}. \quad (3.20)$$

These complex derivatives relate to the second derivatives of the electronic energy with respect to variations of the orbitals (\mathbf{A}_{11} , \mathbf{B}_{11}) and cumulant parameters (\mathbf{A}_{22} , \mathbf{B}_{22}). Similarly, the mixed second derivatives couple variations in the orbitals and cumulant parameters (\mathbf{A}_{12} , \mathbf{B}_{12}). The metric matrix \mathbf{M} has a block-diagonal structure, as a consequence of the linear parametrization of the wavefunction in

eq. (3.5):

$$\mathbf{M} = \begin{pmatrix} \mathbf{S}_{11} & \mathbf{0} & \mathbf{0} & \mathbf{0} \\ \mathbf{0} & \mathbf{1}_2 & \mathbf{0} & \mathbf{0} \\ \mathbf{0} & \mathbf{0} & -\mathbf{S}_{11}^* & \mathbf{0} \\ \mathbf{0} & \mathbf{0} & \mathbf{0} & -\mathbf{1}_2 \end{pmatrix} \quad (3.21)$$

where $\mathbf{1}_2 = \langle \Phi | \mathbf{a}_2^\dagger \mathbf{a}_2 | \Phi \rangle$ is an identity matrix over the space of unique two-body excitations and the orbital metric is defined as follows:

$$\omega \mathbf{S}_{11} = \left. \frac{\partial^2 \{ \langle \Psi(t) | i \dot{\Psi}(t) \rangle \}}{\partial \mathbf{t}_1^\dagger(\omega) \partial \mathbf{t}_1(\omega)} \right|_{f=0} \quad (3.22)$$

Equations for all blocks of \mathbf{E} , \mathbf{M} , and the property gradient vector \mathbf{v}' are shown explicitly in the Supporting Information. We note that, due to the Hermitian nature of the DCT energy functional (3.1), the ODC-12 energy Hessian \mathbf{E} is always symmetric. As a result, in the absence of instabilities (i.e., as long as the Hessian is positive semi-definite), the LR-ODC-12 excitation energies are guaranteed to have real values.

To illustrate the derivation of the LR-ODC-12 energy Hessian, let us consider the diagonal two-body block of \mathbf{E} . Expressing the energy (3.1) using the cumulant expansion (3.2) and differentiating with respect to \mathbf{t}_2 , we obtain:

$$\begin{aligned} \mathbf{A}_{22} = \frac{\partial^2 E}{\partial \mathbf{t}_2^\dagger \partial \mathbf{t}_2} &= f_p^q \frac{\partial^2 \gamma_q^p}{\partial \mathbf{t}_2^\dagger \partial \mathbf{t}_2} + \bar{g}_{pr}^{qs} \frac{\partial \gamma_q^p}{\partial \mathbf{t}_2^\dagger} \frac{\partial \gamma_s^r}{\partial \mathbf{t}_2} \\ &\quad + \frac{1}{4} \bar{g}_{pq}^{rs} \frac{\partial^2 \lambda_{rs}^{pq}}{\partial \mathbf{t}_2^\dagger \partial \mathbf{t}_2} \end{aligned} \quad (3.23)$$

where we have introduced the generalized Fock matrix $f_p^q \equiv h_p^q + \bar{g}_{pr}^{qs} \gamma_s^r$. The derivatives of the one-body density matrix can be expressed in terms of the derivatives of the density cumulant

$$\begin{aligned} \mathbf{A}_{22} = & \mathcal{F}_p^q \frac{\partial^2 \lambda_{qt}^{pt}}{\partial \mathbf{t}_2^\dagger \partial \mathbf{t}_2} + \mathcal{G}_{pr}^{qs} \frac{\partial \lambda_{qt}^{pt}}{\partial \mathbf{t}_2^\dagger} \frac{\partial \lambda_{su}^{ru}}{\partial \mathbf{t}_2} \\ & + \frac{1}{4} \bar{g}_{pq}^{rs} \frac{\partial^2 \lambda_{rs}^{pq}}{\partial \mathbf{t}_2^\dagger \partial \mathbf{t}_2} \end{aligned} \quad (3.24)$$

where the intermediates \mathcal{F}_p^q and \mathcal{G}_{pr}^{qs} can be computed using a transformation of the one- and two-electron integrals to the natural spin-orbital basis (see ?? for details). These cumulant derivatives are straightforward to evaluate from eqs. (3.4) and (3.5) using either algebraic or diagrammatic techniques.

Next, let us outline the derivation of the one-body metric. Substituting eq. (3.5) into eq. (3.22) gives

$$\begin{aligned} \omega \mathbf{S}_{11} = & \frac{1}{2} \frac{\partial^2 \{ \langle \Psi | [i\dot{T}_1^\dagger(t), T_1(t)] | \Psi \rangle \}}{\partial \mathbf{t}_1^\dagger(\omega) \partial \mathbf{t}_1(\omega)} \Big|_{f=0} \\ & - \frac{1}{2} \frac{\partial^2 \{ \langle \Psi | [T_1^\dagger(t), i\dot{T}_1(t)] | \Psi \rangle \}}{\partial \mathbf{t}_1^\dagger(\omega) \partial \mathbf{t}_1(\omega)} \Big|_{f=0} \end{aligned} \quad (3.25)$$

where we have assumed that we are working in the variational orbital basis so that $T_1(t)|_{f=0} = 0$, and $\Psi = \Psi(t)|_{f=0}$ denotes the ground state wavefunction. Using the Fourier expansion of the $\mathbf{t}_1(t)$ parameters (eq. (3.11)), the gradients of the time

derivatives can be evaluated as:

$$\left. \frac{\partial i\dot{T}_1^\dagger(t)}{\partial \mathbf{t}_1^\dagger(\omega)} \right|_{f=0} = -\omega \mathbf{a}_1^\dagger e^{+i\omega t} \quad (3.26)$$

$$\left. \frac{\partial i\dot{T}_1(t)}{\partial \mathbf{t}_1(\omega)} \right|_{f=0} = +\omega \mathbf{a}_1 e^{-i\omega t} \quad (3.27)$$

Substituting eqs. (3.26) and (3.27) into eq. (3.25) and evaluating the gradients of T_1 and T_1^\dagger similarly gives the final working equation for the one-body metric:

$$\begin{aligned} \omega(\mathbf{S}_{11})_{ia,jb} &= \omega \langle \Psi | [a_a^i, a_j^b] | \Psi \rangle \\ &= \omega (\delta_a^b \gamma_j^i - \delta_j^i \gamma_a^b) \end{aligned} \quad (3.28)$$

3.3.4 Linear-Response OLCCD

As we discussed in section 3.3.1, the orbital-optimized linearized coupled cluster doubles method (OLCCD) can be considered as an approximation to the ODC-12 method where all of the non-linear \mathbf{t}_2 terms are neglected. Similarly, we can formulate the linear-response OLCCD method (LR-OLCCD) by linearizing the LR-ODC-12 equations. This simplifies the expressions for the electronic Hessian blocks that involve the second derivatives with respect to \mathbf{t}_2 . For example, for the \mathbf{A}_{22} block, we obtain:

$$\mathbf{A}_{22} = (f_0)_i^j \frac{\partial^2 \lambda_{jr}^{ir}}{\partial \mathbf{t}_2^\dagger \partial \mathbf{t}_2} - (f_0)_a^b \frac{\partial^2 \lambda_{br}^{ar}}{\partial \mathbf{t}_2^\dagger \partial \mathbf{t}_2} + \frac{1}{4} \bar{g}_{pq}^{rs} \frac{\partial^2 \lambda_{rs}^{pq}}{\partial \mathbf{t}_2^\dagger \partial \mathbf{t}_2} \quad (3.29)$$

where $(f_0)_p^q = h_p^q + \bar{g}_{pi}^{qi}$ is the usual (mean-field) Fock operator. Comparing eq. (3.29) with eq. (3.24) from the LR-ODC-12 method, we observe that the former equation can be obtained from the latter by replacing the \mathcal{F}_p^q intermediates with the mean-field Fock matrix elements and ignoring the term that depends on \mathcal{G}_{pr}^{qs} . These simplifications arise from the fact that the \mathcal{F}_p^q and \mathcal{G}_{pr}^{qs} intermediates contain high-order \mathbf{t}_2 contributions that are not included in the linearized LR-OLCCD formulation (see ?? and Ref. 52 for details). For the \mathbf{B}_{22} block, we find that all of the Hessian elements are zero. A complete set of working equations for LR-OLCCD is given in the Supporting Information.

3.4 Computational Details

The LR-ODC-12 and LR-OLCCD methods were implemented as a standalone Python program, which was interfaced with PSI4¹⁶⁷ and PYSCF¹⁶⁸ to obtain the one- and two-electron integrals. To compute excitation energies, our implementation utilizes the multi-root Davidson algorithm,^{169;170} which solves the generalized eigenvalue problem (3.14) by progressively growing an expansion space for the n_{root} lowest generalized eigenvectors of the electronic Hessian and the metric matrix. A key feature of this algorithm is that it avoids storing the Hessian and metric matrices, significantly reducing the amount of memory required by the computations. Our implementation of the energy Hessian was validated by computing the static response function for a dipole perturbation (i.e., the dipole polarizability):

$$\langle\langle \hat{V}; \hat{V} \rangle\rangle_0 = -\mathbf{v}'^\dagger \mathbf{E}^{-1} \mathbf{v}' \quad (3.30)$$

This quantity can also be evaluated numerically as a derivative of the ground state energy

$$\langle\langle\hat{V};\hat{V}\rangle\rangle_0 = \left.\frac{d^2 E}{df^2}\right|_{f=0} \quad (3.31)$$

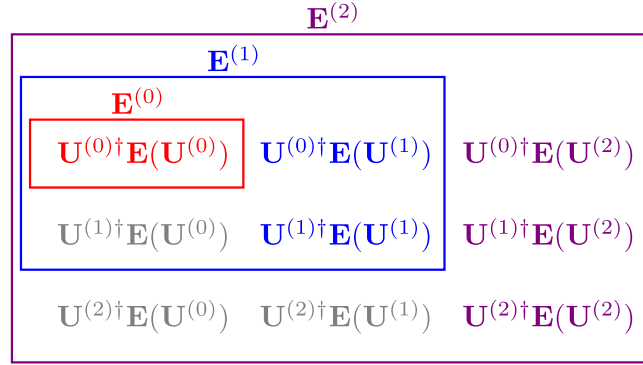
by perturbing the one-electron integrals $h_p^q \leftarrow h_p^q + f\langle\psi_p|\hat{V}|\psi_q\rangle$ and solving the ODC-12 equations for different values of f .

We used Q-CHEM 4.4¹⁷¹ to obtain results from equation-of-motion coupled cluster theory with single and double excitations (EOM-CCSD) and EOM-CCSD with triple excitations in the EOM part [EOM-CC(2,3)]. The MRCC program¹⁷² was used to obtain results for equation-of-motion coupled cluster theory with up to full triple excitations (EOM-CCSDT). All electrons were correlated in all computations. We used tight convergence parameters in all ground-state ($10^{-8} E_h$) and excited-state computations ($10^{-5} E_h$). In ????, the augmented aug-cc-pVTZ and d-aug-cc-pVTZ basis sets of Dunning and co-workers were employed.⁶⁰ For alkenes (section 4.0.1), the ANO-L-pVXZ (X = D, T) basis sets¹⁷³ were used as in Ref. 174. To compute vertical excitation energies in ??, geometries of molecules were optimized using ODC-12 (for LR-ODC-12), OLCCD (for LR-OLCCD), or CCSD [for EOM-CCSD, EOM-CC(2,3), and EOM-CCSDT]. For alkenes in section 4.0.1, the frozen-core MP2/cc-pVQZ geometries were used as in Refs. 174 and 175.

Chapter 4

Linear-Response Density Cumulant Theory for Excited States: Better Algorithms, Bigger Systems

Figure 4.1: A schematic illustrating the sequence of reduced representations of \mathbf{E} in the algorithm employed in this study. The colored entries must be explicitly computed, whereas the gray entries are determined by Hermitian symmetry.



We have implemented the LR-ODC-12 method using a multi-root Davidson algorithm, which solves the linear-response generalized eigenvalue problem by progressively growing an expansion space for the n_{root} lowest generalized eigenvectors of \mathbf{E} and \mathbf{M} . The general procedure looks as follows:

1. Initialize the expansion space with a set of $n_{\text{guess}} \geq n_{\text{root}}$ orthonormal vectors.

$$\mathbf{R}^{(0)} = (\mathbf{u}_1^{(0)} \cdots \mathbf{u}_{n_{\text{guess}}}^{(0)})$$

2. Express the energy-Hessian and metric matrices in the reduced expansion space.

$$\mathbf{E}^{(i)} = \mathbf{R}^{(i)\dagger} \mathbf{E}(\mathbf{R}^{(i)})$$

$$\mathbf{M}^{(i)} = \mathbf{R}^{(i)\dagger} \mathbf{M}(\mathbf{R}^{(i)})$$

3. Solve for the n_{root} highest eigenvectors of the inverse eigenvalue equation.

$$\mathbf{M}^{(i)} \mathbf{z}_k^{(i)} = \mathbf{E}^{(i)} \mathbf{z}_k^{(i)} / \omega_k^{(i)}$$

4. Determine the residual for each root.

$$\mathbf{d}_k^{(i)} = \mathbf{M}(\mathbf{z}_k^{(i)}) - \mathbf{E}(\mathbf{z}_k^{(i)}) / \omega_k^{(i)}$$

If the residual elements are sufficiently small we consider the eigenvectors converged and exit the loop.

5. Form new direction vectors by preconditioning the residuals.

$$\mathbf{g}_k^{(i+1)} = -(\tilde{\mathbf{M}} - \tilde{\mathbf{E}}/\omega_k^{(i)})^{-1} \mathbf{d}_k$$

The tildes in this equation denote diagonal approximations to these matrices.

6. Project out the span of the current expansion space from the new direction vectors.

$$\mathbf{J}^{(i+1)} = (\mathbf{1} - \mathbf{R}^{(i)} \mathbf{R}^{(i)\dagger})(\mathbf{g}_1^{(i+1)} \dots \mathbf{g}_{n_{\text{root}}}^{(i+1)})$$

7. Determine an orthonormal basis for the new direction vectors using a compressed singular value decomposition, and add these new vectors to the expansion space.

$$\mathbf{J}^{(i+1)} \approx \mathbf{U}^{(i+1)} \mathbf{\Sigma}^{(i+1)} \mathbf{V}^{(i+1)\dagger}$$

$$\mathbf{R}^{(i+1)} \leftarrow (\mathbf{R}^{(i)} \quad \mathbf{U}^{(i+1)})$$

8. Increment i and return to step 2.

A key feature of this algorithm is that we never need to explicitly construct the energy-Hessian and metric matrices in memory, only their images over the expansion space.

For the diagonal approximations in step 5 we use the following.

$$\tilde{\mathbf{S}}_{11} \equiv \mathbf{1}_1$$

$$(\tilde{\mathbf{A}}_{11})_{ia,ia} \equiv -f_i^i + f_a^a$$

$$(\tilde{\mathbf{A}}_{22})_{ijab,ijab} \equiv -\mathcal{F}_i^i - \mathcal{F}_j^j - \mathcal{F}_a^a - \mathcal{F}_b^b$$

These can also be used to construct the initial expansion space in step 1, namely by including unit vectors for the n_{guess} smallest positive entries in $\tilde{\mathbf{E}}$.

4.0.1 Ethylene, Butadiene, and Hexatriene

Table 4.1: Vertical excitation energies computed using LR-OLCCD, LR-ODC-12, and EOM-CCSD for the low-lying electronic states of ethylene (C_2H_4), butadiene (C_4H_6), and hexatriene (C_6H_8). Computations employed the ANO-L-pVDZ (for C_4H_6 and C_6H_8) and ANO-L-pVTZ (for C_2H_4) basis sets and the MP2/cc-pVQZ optimized geometries. For LR-OLCCD and LR-ODC-12, oscillator strengths of the allowed transitions are given in parentheses. All electrons were correlated in all computations. Also shown are the excitation energies from the frozen-core semistochastic heat-bath CI (SHCI) method, extrapolated to full CI limit.

		EOM-CCSD	LR-OLCCD	LR-ODC-12	SHCI ^a
C_2H_4	$^3\text{B}_{1\text{u}}$	4.46	4.66	4.52	4.59
	$^1\text{B}_{1\text{u}}$	8.14	8.20 (1.8)	8.14 (1.9)	8.05
C_4H_6	$^3\text{B}_{\text{u}}$	3.20	3.58	3.43	3.37
	$^1\text{B}_{\text{u}}$	6.53	6.76 (4.2)	6.67 (4.4)	6.45
	$^1\text{A}_{\text{g}}$	7.28	7.14	6.81	6.58
C_6H_8	$^3\text{B}_{\text{u}}$	2.64	3.01	2.83	2.77
	$^1\text{B}_{\text{u}}$	5.60	5.89 (6.5)	5.74 (8.1)	5.59
	$^1\text{A}_{\text{g}}$	6.55		5.73	5.58

^a The SHCI computations used the same basis sets and optimized geometries as those used for LR-OLCCD, LR-ODC-12, and EOM-CCSD.

Chapter 5

Conclusion

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Appendix A

Extended Results

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