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Assessment of the Validity of Some Minnesota Density Functionals for the Prediction of the Chemical Reactivity of the SYBR Green I and Ethidium Bromide Nucleic Acid Stains

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Additional information is available at the end of the chapter

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Abstract

This research work has assessed many Minnesota density functionals to find their molecular structure and electronic properties possessed by SYBR green I (SYBRGI) and ethidium bromide (EtBr) nucleic acid stains. In the determination of the global descriptors that come up from conceptual density functional theory (CDFT), the processes include: Self-Consistent Field Energy Differences (Δ SCF) and higher occupied molecular orbital (HOMO) and lower unoccupied molecular orbital (LUMO) frontier orbitals energies. Regarding the deduced outcomes for the conceptual DFT indices, many of the descriptors have been adjusted to achieve the “Koopmans in DFT (KID)” process. It has also been shown that the only density functionals that confirm this approximation are the range-separated hybrids (RSH).

Keywords: computational chemistry, SYBR green I, chemical reactivity theory, molecular modeling, conceptual DFT, ethidium bromide

1. Introduction

The chemical reactivity theory [also known as the conceptual density functional theory (CDFT)] is a vital technique that is used to predict, evaluate, and interpret the results from chemical processes [1–4].

Research done by Parr and his associates [1] reveals that several theories and models have been discovered after the evaluation of the molecular system with the use of DFT. Almost all the discovered theories are helpful in research because they enable scholars to achieve quantitative forecasts of a chemical reactivity system. In addition to this, the theories can further be quantified and are generally termed as conceptual DFT descriptors.

To obtain the quantitative figures of conceptual descriptors, it may be necessary to analyze the Kohn-Sham theory by calculating the energy system, molecular density, and the orbital energies associated with frontier orbitals [i.e., higher occupied molecular orbital (HOMO) and lower unoccupied molecular orbital (LUMO)] [5–10].

For research on a molecular system, the first activity before proceeding is selection of the model chemistry. Model chemistry is the collection of basic set, density function, together with an implied solvent model that is known to be consistent for the problem under research. Several studies provide insights on the way to choose the model chemistry. A researcher may also decide to preview past studies when choosing the model.

Even though the fundamentals of DFT reveal that universal functional density is present and that computations using this function can be used to obtain all the features of the system, it is always necessary in practical cases that one refers to the estimated density functionals that have been established for the past 3 decades. For the approximate functionals, almost all of them are perfectly fit to be used in estimating some features, while some can be used for estimating other features. In separate scenarios, you can encounter density functionals that are perfectly fit for estimating the features of a given molecular system and a functional group. It is also important to assess separate density functionals for a separate functional group which can be added to the molecular system under research.

When researching on chemical reactivity (which is a process that entails the transfer of electrons), a person performs computations for both ground and open systems, i.e., cation and anion. It is not easy to obtain consistent outcomes using these computations (when diffuse functions should be a part of the basis set) [5–10]. This necessitates adoption of a more consistent technique that provides all the data that a person will require directly from the outcomes of the computations at ground state in the molecular system under research. In addition to this, a person may also want to find the deionization ability together with the electron affinity of any system being researched without having to calculate the radical cation and anion. This can be determined by the Koopmans' theorem [7–10] that relies on Hartree-Fock Theory, which states that the energy of the HOMO (i.e., $I = -\epsilon_H$) can be used to estimate the ionization potential. Alternatively, the electron affinity can be estimated using the minus the energy of the LUMO (i.e., $A = -\epsilon_L$).

The legitimacy of the Koopmans' theorem is yet again a contentious issue because of the existing difference between the fundamental band gap and the HOMO and the LUMO gaps. This can be termed as derivative discontinuity. It has again been discovered that an exact physical description may be assigned to Kohn-Sham HOMO using "the Kohn-Sham analogue of Koopmans' theorem in Hartree-Fock theory" (this theory explains that in the exact theory, the KS HOMO is opposite and same as the ionization potential) [11–14]. The effects brought about by the difference between the fundamental band gap and the HOMO and the LUMO gaps have ensured that no Koopmans' theorem creates a direct relationship between the LUMO energy and the electron affinity. To eliminate these effects, a suggestion has been made by scholars to conceive that the ionization potential of the $N + 1$ electron system (anion) is almost equal to the electron affinity of N electron system [15]. Regarding the range-separated hybrids (RSH) density functionals [16–18], e.g., that the repulsive coulomb potential has to be separated in the long-range (LR) and short-range (SR) terms, e.g., via $r^{-1} = r^{-1} \operatorname{erf}(\gamma r) + r^{-1} \operatorname{erfc}(\gamma r)$, with γ representing the range-separation parameter, it was highlighted by Kronik et al.

[15] that the legitimacy of the Koopmans' theorem may be approximately approved with a well-informed choice of this final parameter.

Use of the γ -tuning technique can be useful in upgrading the features projected by the density functionals. This is due to better utilization of the of Koopmans' theorem, leading to a superior understanding of frontier orbitals energies with I and A. An example can be cited from Lima et al. [19], who just demonstrated a better explanation of the optical features of carotenoids through tuning of some LR density functionals that are linked.

This therefore illustrates that conformity of any given density functional can be determined by assessing how it has adopted the "Koopmans' in DFT" (KID) process, which guides its behavior to be almost equal to the ideal density functional. This is essential for any precise computation of the conceptual DFT descriptors that help in forecasting and analysis of chemical reactivity in molecular systems. Still, the γ -tuning technique for range-separated hybrids density functional is system dependent. This implies that separate density functionals are to be used in the computations of the descriptors for separate molecular features. We are then going to concentrate on part of the density functionals that have displayed the required precision in physics and chemistry [20].

The main aim of this study is to do a comparative research relating to the performance of the just identified Minnesota family of density functionals for the account of the chemical reactivity of two nucleic acids intercalating stains, SYBR green I (SYBRGI) [21] and ethidium bromide (EtBr) [22]. The molecular structures of the two are shown in **Figure 1**.

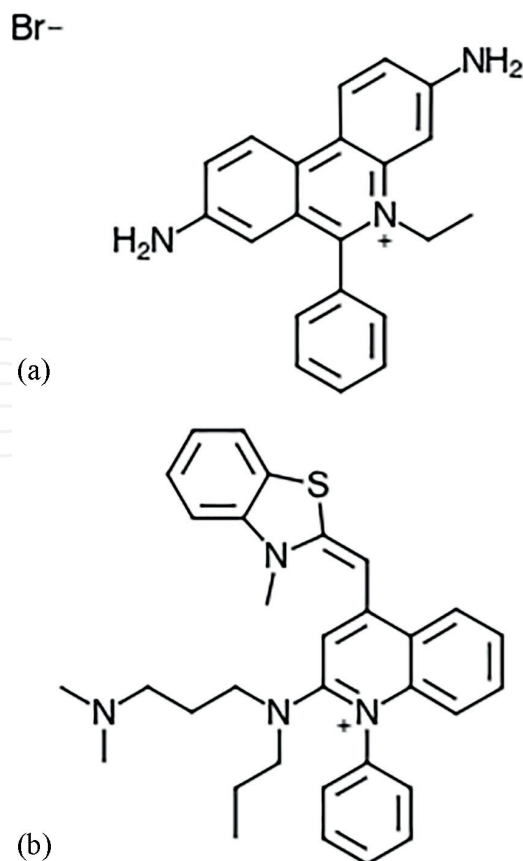


Figure 1. Molecular structures of (a) ethidium bromide (EtBr) and (b) SYBR green I (SYBRGI).

2. Theoretical background

Within the context of DFT [2, 23], the chemical potential μ , which estimates the escaping tendency of the electron from the equilibrium point is stated as follows:

$$\mu = \left(\frac{\partial E}{\partial N} \right)_{v(r)} \quad (1)$$

where χ represents the electronegativity.

Chemical hardness is represented by η , which is the opposition to charge transfer:

$$\eta = \left(\frac{\partial^2 E}{\partial N^2} \right)_{v(r)} \quad (2)$$

Employing a finite difference approximation and the Koopmans's theorem [7–10], we can write the above equation as:

$$\mu = -\frac{1}{2}(I + A) \approx \frac{1}{2}(\varepsilon_L + \varepsilon_H) = -\chi_K \quad (3)$$

$$\eta = (I - A) \approx (\varepsilon_L - \varepsilon_H) = \eta_K \quad (4)$$

where ε_H is the HOMO energy and ε_L is the LUMO energy.

An expression for the electrophilicity index ω is as below:

$$\omega = \frac{\mu^2}{2\eta} = \frac{(I + A)^2}{4(I - A)} \approx \frac{(\varepsilon_L + \varepsilon_H)^2}{4(\varepsilon_L - \varepsilon_H)} = \omega_K \quad (5)$$

Expressions for electrodonating ω^- and electroaccepting ω^+ powers are as below [24]:

$$\omega^- = \frac{(3I + A)^2}{4(I - A)} \approx \frac{(3\varepsilon_H + \varepsilon_L)^2}{16\eta_K} = \omega_K^- \quad (6)$$

and

$$\omega^+ = \frac{(I + 3A)^2}{4(I - A)} \approx \frac{(\varepsilon_H + 3\varepsilon_L)^2}{16\eta_K} = \omega_K^+ \quad (7)$$

To obtain a comparison for ω^+ and $-\omega^-$, the explanation below for net electrophilicity has been suggested [25]:

$$\Delta\omega^\pm = \omega^+ - (-\omega^-) = \omega^+ + \omega^- \approx \omega_K^+ - (-\omega_K^-) = \omega_K^+ + \omega_K^- = \Delta\omega_K^\pm \quad (8)$$

3. Settings and computational details

In this research project, each of the computations was done using Gaussian 09 programs [26] and the density functional methods as compelled in the computational package. The gradient method was used to obtain the equilibrium geometries of molecules in this research. Additionally, vibration frequencies and the force parameters were estimated through computation of analytical frequencies on still areas after optimization to check whether they were the actual minima. Def2SVP was used in this research project as the basic set for optimization of geometry and frequencies. Computation of the electronic features was achieved using Def2TZVP basic set [27, 28].

We chose a group of Minnesota density functionals, which give consistent outcomes when computing the molecular structures and systems during the research activities. The group below were selected: M11, which falls under RSH meta-generalized gradient approximation (GGA) [29]; M11L, which falls under double-range local meta-GGA [30]; MN12L, which falls under nonseparable meta-nonseparable gradient approximation (NGA) [31]; MN12SX, which falls under nonseparable hybrid nonseparable meta-NGA [32]; N12, which falls under nonseparable gradient estimation [33]; N12SX, which falls under RSH nonseparable gradient estimation [32]; SOGGA11, which falls under generalized gradient approximation (GGA) density functional [34]; and SOGGA11X, which falls under generalized gradient approximation density functional [35]. GGA can be explained as a functional whereby the functional relies on the both the up down spin densities and the minimized gradient. Nonseparable gradient approximation (NGA) can be explained as a functional that relies on up down spin densities and minimized gradient and falls under nonseparable. In the current research, each of the computations was done where the solvent used was water and by doing the calculations in conformity to the Solvation Model based on Density (SMD) solvation standard [36].

4. Results and discussion

Firstly, the molecular structures of SYBRGI and EtBr were first optimized by MOL structures, then by finding the most stable conformers through Avogadro program [37, 38]. This was done through random sampling with molecular mechanic techniques. After the optimization, the resulting conformers were then re-optimized with MN12SX, M11L, M11, N12, SOGGA11X, and SOGGA11 density functionals. In addition, Def2SVP basic set and SMD solvation model were used, water being used as a solvent.

The HOMO and LUMO energies (in eV), the ionization potential I and electron affinity A (in eV), electronegativity χ , chemical hardness η , global electrophilicity ω , electrodonating ω^- , and electroaccepting ω^+ powers, and the net electrophilicity $\Delta\omega^\pm$ of the EtBr and SYBRGI molecules calculated with the same density functionals and solvation model are presented in **Tables 1** and **2**, respectively. The upper part of the tables shows the results derived assuming the validity of the Kid procedure (hence the subscript K), and the lower part of the tables shows the results derived from the calculated vertical I and A through a Δ SCF technique.

Property	M11	M11L	MN12L	MN12SX	N12	N12SX	SOGGA11	SOGGA11X
HOMO	-7.535	-4.956	-4.684	-5.187	-4.164	-4.951	-4.151	-6.108
LUMO	-0.535	-3.028	-2.613	-2.576	-2.674	-2.511	-3.121	-1.808
χ_K	4.035	3.992	3.649	3.882	3.419	3.731	3.636	3.958
η_K	7.000	1.928	2.071	2.611	1.490	2.440	1.030	4.300
ω_K	1.163	4.133	3.213	2.886	3.924	2.852	6.420	1.822
ω_K^-	4.781	10.383	8.380	7.876	9.651	7.723	14.722	5.891
ω_K^+	0.746	6.391	4.732	3.994	6.232	3.992	11.086	1.933
$\Delta\omega_K^\pm$	5.527	16.774	13.112	11.869	15.883	11.714	25.808	7.824
I	5.585	5.183	4.883	5.216	4.520	4.970	4.819	5.385
A	2.711	2.782	2.408	2.624	2.344	2.579	2.788	2.663
χ	4.148	3.983	3.646	3.920	3.432	3.775	3.804	4.024
η	2.874	2.401	2.475	2.592	2.177	2.392	2.031	2.721
ω	2.994	3.303	2.685	2.965	2.706	2.978	3.562	2.975
ω^-	8.241	8.748	7.348	8.051	7.263	7.993	9.152	8.132
ω^+	4.093	4.786	3.702	4.131	3.831	4.219	5.348	4.108
$\Delta\omega^\pm$	12.335	13.514	11.050	12.182	11.094	12.212	14.500	12.241

Table 1. HOMO and LUMO energies (in eV), the ionization potential I and electron affinity A (in eV), electronegativity χ , chemical hardness η , global electrophilicity ω , electrodonating ω^- and electroaccepting ω^+ powers, and the net electrophilicity $\Delta\omega^\pm$ of the EtBr molecule.

For examining the outcomes to determine if the KID process is fulfilled, and the drive from past works [15, 19], we have come up with descriptors having the ability to compare the outcomes from HOMO and LUMO computations with those attained using vertical I and A and a Δ SCF technique. It should again be known that we have no plans to form a gap fitting by reducing the descriptor. We plan to determine if the density functionals employed in this research contain the fixed range parameter γ that helps in effective execution of the KID process. It is somehow astonishing that our research at present lacks the parameter γ . We also included a minus of the energy of the LUMO of the neutral system instead of using A as minus of HOMO of the electron system [15, 19].

The initial three descriptors are associated with the basic accomplishment of "Koopmans in DFT" estimation by associating ε_H with $-I$, ε_L with $-A$, and their responses in explaining the HOMO-LUMO gap:

$$J_I = |\varepsilon_H + E_{gs}(N-1) - E_{gs}(N)| \quad (9)$$

$$J_A = |\varepsilon_L + E_{gs}(N+1) - E_{gs}(N)| \quad (10)$$

$$J_{HL} = \sqrt{J_I^2 + J_A^2} \quad (11)$$

Property	M11	M11L	MN12L	MN12SX	N12	N12SX	SOGGA11	SOGGA11X
HOMO	-7.593	-5.132	-4.924	-5.325	-4.587	-5.160	-4.779	-6.099
LUMO	-0.558	-2.933	-2.569	-2.545	-2.618	-2.460	-2.959	-1.768
χ_K	4.075	4.033	3.747	3.935	3.603	3.810	3.869	3.933
η_K	7.034	2.199	2.355	2.780	1.970	2.700	1.820	4.331
ω_K	1.181	3.698	2.980	2.785	3.295	2.688	4.112	1.786
ω^-_K	4.839	9.550	7.981	7.711	8.514	7.449	10.273	5.809
ω^+_K	0.763	5.517	4.234	3.776	4.911	3.639	6.404	1.876
$\Delta\omega^\pm_K$	5.602	15.067	12.214	11.486	13.425	11.088	16.677	7.685
I	5.407	5.302	5.062	5.263	4.767	5.074	4.980	5.252
A	2.643	2.747	2.400	2.589	2.358	2.521	2.690	2.564
χ	4.025	4.024	3.731	3.926	3.563	3.797	3.835	3.908
η	2.764	2.555	2.661	2.674	2.409	2.553	2.290	2.688
ω	2.931	3.169	2.615	2.882	2.635	2.824	3.210	2.841
ω^-	8.048	8.509	7.262	7.894	7.202	7.706	8.481	7.803
ω^+	4.023	4.485	3.531	3.968	3.639	3.909	4.646	3.895
$\Delta\omega^\pm$	12.071	12.994	10.972	11.862	10.841	11.615	13.128	11.699

Table 2. HOMO and LUMO energies (in eV), the ionization potential I and electron affinity A (in eV), electronegativity χ , chemical hardness η , global electrophilicity ω , electrodonating ω^- and electroaccepting ω^+ powers, and the net electrophilicity $\Delta\omega^\pm$ of the SYBRGI molecule.

Four separate descriptors will then be used to examine how the density functionals under research will help in forecasting the electronegativity χ , the chemical hardness η , the global electrophilicity ω , and the collection of conceptual DFT descriptors through deliberation of the energies of the HOMO and LUMO or the vertical I and A:

$$J_\chi = |\chi - \chi_K| \quad (12)$$

$$J_\eta = |\eta - \eta_K| \quad (13)$$

$$J_\omega = |\omega - \omega_K| \quad (14)$$

$$J_{D1} = \sqrt{J_\chi^2 + J_\eta^2 + J_\omega^2} \quad (15)$$

D1 represents the initial collection of conceptual DFT descriptors.

Finally, we came up with four extra descriptors to determine the success of the density functionals under research in forecasting of electrodonating power ω^- , the electroaccepting power ω^+ , the net electrophilicity $\Delta\omega^\pm$, together with the four descriptors combined and considering the HOMO and LUMO energies or the vertical I and A:

$$J_{\omega^-} = |\omega^- - \omega_K^-| \quad (16)$$

$$J_{\omega^+} = |\omega^+ - \omega_K^+| \quad (17)$$

$$J_{\Delta\omega\pm} = |\Delta\omega \pm -\Delta\omega\pm_K| \quad (18)$$

$$J_{D2} = \sqrt{J_{\omega^-}^2 + J_{\omega^+}^2 + J_{\Delta\omega\pm}^2} \quad (19)$$

D2 represents the second collection of conceptual DFT descriptors.

The results of the calculations of $J_I, J_A, J_{HL}, J_X, J_\eta, J_\omega, J_{D1}, J_{\omega^-}, J_{\omega^+}, J_{\Delta\omega\pm}$, and J_{D2} for the EtBr and SYBRGI are displayed in **Tables 3** and **4**, respectively.

As shown in **Tables 1** and **2**, and the outcomes from **Tables 3** and **4**, the precision provided by the KID process is outstanding for the MN12SX, which falls under RSH meta-NGA, and N12SX, which falls under RSH NGA density functionals. In reality, values for J_I, J_A and J_{HL} is not zero. However, the values found can satisfactorily be likened to the past studies of Lima et al. [19], whereby the minima were found by selecting a parameter that imposes such a trend.

The outcomes are necessary because they reveal that we should not depend on J_I, J_A , and J_{HL} alone, i.e., if we depend on outcomes from J_X , alone, almost all the values will near zero. For the remaining descriptors, only MN12SX and N12SX reveal such trends. This shows that outcomes for J_X can be due to elimination of errors.

Authentication of the KID process is not done correctly by the GGA (SOGGA11) and hybrid-GGA (SOGGA11X). Local density functionals like M11L, MN12L, and N12 are also inappropriate.

Descriptor	M11	M11L	MN12L	MN12SX	N12	N12SX	SOGGA11	SOGGA11X
J_I	1.950	0.228	0.199	0.028	0.356	0.020	0.668	0.724
J_A	2.176	0.246	0.205	0.048	0.331	0.068	0.333	0.855
J_{HL}	2.922	0.335	0.285	0.055	0.486	0.071	0.747	1.120
J_X	0.113	0.009	0.003	0.038	0.013	0.044	0.168	0.066
J_η	4.127	0.473	0.403	0.019	0.687	0.048	1.001	1.579
J_ω	1.831	0.830	0.528	0.079	1.219	0.126	2.858	1.154
J_{D1}	4.516	0.955	0.665	0.090	1.399	0.142	3.033	1.957
J_{ω^-}	3.460	1.635	1.033	0.175	2.388	0.271	5.570	2.242
J_{ω^+}	3.347	1.626	1.030	0.137	2.401	0.227	5.738	2.176
$J_{\Delta\omega^+}$	6.808	3.260	2.062	0.313	4.788	0.498	11.307	4.417
J_{D2}	8.338	3.993	2.526	0.384	5.864	0.611	13.849	5.410

Table 3. Descriptors $J_I, J_A, J_{HL}, J_X, J_\eta, J_\omega, J_{D1}, J_{\omega^-}, J_{\omega^+}, J_{\Delta\omega\pm}$, and J_{D2} for the ethidium bromide (EtBr) molecule calculated from the results of **Table 1**.

Descriptor	M11	M11L	MN12L	MN12SX	N12	N12SX	SOGGA11	SOGGA11X
J_1	2.185	0.170	0.137	0.062	0.180	0.086	0.201	0.847
J_A	2.085	0.187	0.169	0.044	0.259	0.061	0.269	0.796
J_{HL}	3.020	0.252	0.218	0.076	0.316	0.105	0.336	1.162
J_X	0.050	0.009	0.016	0.009	0.040	0.012	0.034	0.025
J_η	4.270	0.356	0.306	0.106	0.439	0.147	0.470	1.643
J_ω	1.751	0.529	0.365	0.097	0.660	0.136	0.902	1.055
J_{D1}	4.615	0.638	0.477	0.144	0.794	0.201	1.018	1.952
$J_{\omega-}$	3.209	1.041	0.719	0.183	1.312	0.257	1.792	1.994
$J_{\omega+}$	3.259	1.032	0.703	0.192	1.272	0.270	1.758	2.019
$J_{\Delta\omega+}$	6.469	2.073	1.422	0.376	2.584	0.527	3.550	4.013
J_{D2}	7.923	2.539	1.742	0.460	3.165	0.646	4.348	4.915

Table 4. Descriptors $J_1, J_A, J_{HL}, J_X, J_\eta, J_\omega, J_{D1}, J_{\omega-}, J_{\omega+}, J_{\Delta\omega+}$ and J_{D2} for the SYBR green I (SYBRGI) molecule calculated from the results of **Table 2**.

It is vital to know that even though the RSH hybrid NGA and RSH meta-NGA density functionalities are necessary when computing the conceptual DFT descriptors, it is a different case for RSH GGA (M11) density functional. According to **Tables 1** and **2**, this functional doesnot provide enough explanation concerning LUMO energy, and this can be due to an inaccurate figure of γ in the functional. A fine tuning of γ can handle the issue.

5. Conclusions

Weighing on the outcomes from this research work, DFT-based reactivity descriptors like electronegativity, chemical hardness, global electrophilicity, electrodonating, and electro-accepting powers, and net electrophilicity can be used to forecast EtBr's chemical reactivity.

It has also been illustrated that the KID process can effectively be implemented by the RSH meta-NGA (MN12SX) and the RSH NGA (N12SX) density functionalities. They can then be used in place of the tuned density functionals using a gap-fitting process, and we believe that such a trend can be helpful when analyzing the chemical reactivity of bigger molecular systems.

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References

- [1] Parr RG, Yang W. Density Functional Theory of Atoms and Molecules. New York: Oxford University Press; 1989
- [2] Geerlings P, De Proft F, Langenaeker W. Conceptual density functional theory. *Chemical Reviews*. 2003;**103**:1793-1873
- [3] Toro-Labbé A, editor. Theoretical Aspects of Chemical Reactivity. Vol. 19. Amsterdam: Elsevier Science; 2007
- [4] Chattaraj PK, editor. Chemical Reactivity Theory – A Density Functional View. Boca Raton: CRC Press, Taylor & Francis Group; 2009
- [5] Huzinaga S, Andzelm J, Klobukowski M, Radzio-Audzelm E, Sakai Y, Tatewaki H, Gaussian Basis Sets for Molecular Calculations. Amsterdam: Elsevier; 1984
- [6] Easton R, Giesen D, Welch A, Cramer C, Truhlar D. The MIDI! basis set for quantum mechanical calculations of molecular geometries and partial charges. *Theoretical Chemistry Accounts*. 1996;**93**:281-301
- [7] Lewars E. Computational Chemistry – Introduction to the Theory and Applications of Molecular and Quantum Mechanics. Dordrecht: Kluwer Academic Publishers; 2003
- [8] Young D. Computational Chemistry – A Practical Guide for Applying Techniques to Real-World Problems. New York: John Wiley & Sons; 2001
- [9] Jensen F. Introduction to Computational Chemistry. 2nd ed. Chichester, England: John Wiley & Sons; 2007
- [10] Cramer C. Essentials of Computational Chemistry – Theories and Models. 2nd ed. Chichester, England: John Wiley & Sons; 2004

- [11] Perdew JP, Parr RG, Levy M, Balduz JLJ. Density-functional theory for fractional particle number: Derivative discontinuities of the energy. *Physical Review Letters*. 1982;**49**: 1691-1694
- [12] Almbladh CO, von Barth U. Exact results for the charge and spin densities, exchange-correlation potentials, and density-functional eigenvalues. *Physical Review B*. 1985;**31**: 3231-3244
- [13] Perdew JP, Burke K, Ernserhof M. Erratum: Generalized gradient approximation made simple. *Physical Review Letters*. 1997;**78**:1396
- [14] Levy M, Perdew JP, Sahni V. Exact differential equation for the density and ionization energy of a many-particle system. *Physical Review A*. 1984;**30**:2745-2748
- [15] Kronik L, Stein T, Refaely-Abramson S, Baer R. Excitation gaps of finite-sized systems from optimally tuned range-separated hybrid functionals. *Journal of Chemical Theory and Computation*. 2012;**8**(5):1515-1531
- [16] Savin A. Beyond the Kohn-Sham determinant. In: *Recent Advances in Density Functional Methods*. Singapore: World Scientific; 2011. Ch. 4. pp. 129-153
- [17] Leininger T, Stoll H, Werner HJ, Savin A. Combining long-range configuration interaction with short-range density functionals. *Chemical Physics Letters*. 1997;**275**(3):151-160
- [18] Savin A, Flad HJ. Density functionals for the Yukawa electron-electron interaction. *International Journal of Quantum Chemistry*. 1995;**56**(4):327-332
- [19] Lima IT, Prado ADS, Martins JBL, de Oliveira Neto PH, Ceschin AM, da Cunha WF, da Silva Filho DA. Improving the description of the optical properties of carotenoids by tuning the long-range corrected functionals. *The Journal of Physical Chemistry A*. 2016;**120**(27):4944-4950
- [20] Peverati R, Truhlar DG. Quest for a universal density functional: The accuracy of density functionals across a broad spectrum of databases in chemistry and physics. *Philosophical Transactions. Series A, Mathematical, Physical, and Engineering Sciences*. 2014;**372**(2011): 20120476
- [21] Green F, editor. *The Sigma-Aldrich Handbook of Stains, Dyes, and Indicators*. Milwaukee, Wis: Aldrich Chemical Co.; 1990
- [22] Sabnis R, editor. *Handbook of Biological Dyes and Stains*. New Jersey, Hoboken: John Wiley & Sons; 2010
- [23] Parr RG, Yang W. Density functional approach to the frontier-electron theory of chemical reactivity. *Journal of the American Chemical Society*. 1984;**106**:4049-4050
- [24] Gázquez JL, Cedillo A, Vela A. Electrodonating and electroaccepting powers. *Journal of Physical Chemistry A*. 2007;**111**(10):1966-1970
- [25] Chattaraj PK, Chakraborty A, Giri S. Net electrophilicity. *Journal of Physical Chemistry A*. 2009;**113**(37):10068-10074

- [26] Frisch MJ, Trucks GW, Schlegel HB, Scuseria GE, Robb MA, Cheeseman JR, Scalmani G, Barone V, Mennucci B, Petersson GA, Nakatsuji H, Caricato M, Li X, Hratchian HP, Izmaylov AF, Bloino J, Zheng G, Sonnenberg JL, Hada M, Ehara M, Toyota K, Fukuda R, Hasegawa J, Ishida M, Nakajima T, Honda Y, Kitao O, Nakai H, Vreven T, Montgomery Jr JA, Peralta JE, Ogliaro F, Bearpark M, Heyd JJ, Brothers E, Kudin KN, Staroverov VN, Kobayashi R, Normand J, Raghavachari K, Rendell A, Burant JC, Iyengar SS, Tomasi J, Cossi M, Rega N, Millam JM, Klene M, Knox JE, Cross JB, Bakken V, Adamo C, Jaramillo J, Gomperts R, Stratmann RE, Yazyev O, Austin AJ, Cammi R, Pomelli C, Ochterski JW, Martin RL, Morokuma K, Zakrzewski VG, Voth GA, Salvador P, Dannenberg JJ, Dapprich S, Daniels AD, Farkas O, Foresman JB, Ortiz JV, Cioslowski J, Fox DJ. Gaussian 09 Revision D.01. Wallingford CT: Gaussian Inc.; 2016
- [27] Weigend F, Ahlrichs R. Balanced basis sets of split valence, triple zeta valence and quadruple zeta valence quality for H to Rn: Design and assessment of accuracy. *Physical Chemistry Chemical Physics*. 2005;7:3297-3305
- [28] Weigend F. Accurate coulomb-fitting basis sets for H to R. *Physical Chemistry Chemical Physics*. 2006;8:1057-1065
- [29] Peverati R, Truhlar DG. Improving the accuracy of hybrid meta-GGA density functionals by range separation. *The Journal of Physical Chemistry Letters*. 2011;2(21):2810-2817
- [30] Peverati R, Truhlar DG. M11-L: A local density functional that provides improved accuracy for electronic structure calculations in chemistry and physics. *The Journal of Physical Chemistry Letters*. 2012;3(1):117-124
- [31] Peverati R, Truhlar DG. An improved and broadly accurate local approximation to the exchange-correlation density functional: The MN12-L functional for electronic structure calculations in chemistry and physics. *Physical Chemistry Chemical Physics*. 2012;14(38):13171-13174
- [32] Peverati R, Truhlar DG. Screened-exchange density functionals with broad accuracy for chemistry and solid-state physics. *Physical Chemistry Chemical Physics*. 2012;14(47):16187-16191
- [33] Peverati R, Truhlar DG. Exchange-correlation functional with good accuracy for both structural and energetic properties while depending only on the density and its gradient. *Journal of Chemical Theory and Computation*. 2012;8(7):2310-2319
- [34] Peverati R, Zhao Y, Truhlar DG. Generalized gradient approximation that recovers the second-order density-gradient expansion with optimized across-the-board performance. *The Journal of Physical Chemistry Letters*. 2011;2(16):1991-1997
- [35] Peverati R, Truhlar DG. Communication: A global hybrid generalized gradient approximation to the exchange-correlation functional that satisfies the second-order density-gradient constraint and has broad applicability in chemistry. *The Journal of Chemical Physics*. 2011;135(19):191102

- [36] Marenich A, Cramer C, Truhlar DG. Universal solvation model based on solute electron density and a continuum model of the solvent defined by the bulk dielectric constant and atomic surface tensions. *Journal of Physical Chemistry B*. 2009;**113**:6378-6396
- [37] Avogadro: An Open-Source Molecular Builder and Visualization Tool – Version 1.2.0. 2016. Available from: <http://avogadro.openmolecules.net>
- [38] Hanweel M, Curtis DE, Lonie D, Vandermeersch T, Zurek E, Hutchison G. Avogadro: An advanced semantic chemical editor, visualization, and analysis platform. *Journal of Cheminformatics*. 2012;**4**:17

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