Density-Dependent Onset of the Long-Range Exchange: A Key to Donor-Acceptor Properties

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Abstract

Quantum mechanical methods based on the density functional theory (DFT) offer a realistic possibility of first-principles design of organic donor-acceptor systems and engineered band-gap materials. This promise is contingent upon the ability of DFT to predict one-particle states accurately. Unfortunately, approximate functionals fail to align the orbital energies with ionization potentials. We describe a new paradigm for achieving this alignment. In the proposed model, an average electron—exchange hole separation controls the onset of the orbital-dependent exchange in approximate range-separated functionals. The correct description of one-particle states is thus achieved.
without explicit electron removal or attachment. Extensive numerical tests show that
the proposed method provides physically sound orbital gaps and leads to excellent
predictions of charge-transfer excitations and other properties critically depending on
the tail of the electron density.

Introduction

A nonlocal exchange functional is critical to the description of properties that depend on
the tail of the electron density, such as charge-transfer (CT) and Rydberg excitations,\(^1\)\(^-\)\(^3\)
repulsive part of the van der Waals potential,\(^4\) and frontier orbitals.\(^5\) In range-separated
(RS) functionals,\(^6\)\(^-\)\(^8\) a nonlocal, orbital-dependent exchange functional is introduced in the
long-range part of the electron-electron interaction. The critical issue is at what distance
the long range should commence. This paper presents a model that explains the underlying
physics of a molecular system behind the proper onset of the long-range exchange. An
efficient and reliable functional is derived for the properties depending on the density tail.

A mixture of a local density functional approximation (DFA) and orbital-dependent
Hartree-Fock (HF) exchange was first proposed by Becke\(^9\) and subsequently incorporated
into a variety of models. The non-local, orbital-dependent exchange absorbs a part of the
missing discontinuity of the exchange-correlation potential\(^10\) plaguing local DFAs.\(^11\)\(^,\)\(^12\) Al-
though hybrid DFAs improve on the quality of molecular properties, several issues remain.
Most notably, qualitative errors in donor-acceptor properties, fundamental gaps from or-
bital energies, and CT excitations were attributed to the persistence of the many-electron
self-interaction error.\(^13\)\(^,\)\(^14\) These deficiencies of hybrid DFAs can be alleviated by applying
different models for short- and long-range electronic interaction according to the decompo-
sition of the electron-electron interaction operator:\(^6\)\(^-\)\(^8\)
\[
\frac{1}{r_{12}} = \frac{\text{erfc}(\omega r_{12})}{r_{12}} + \frac{\text{erf}(\omega r_{12})}{r_{12}}. \tag{1}
\]
The decomposition is followed by the separation of short- and long-range parts out of the total exchange hole,

\[ h_X^\sigma(r_1, r_2) = h_{X,SR}^\sigma(r_1, r_2) + h_{X,LR}^\sigma(r_1, r_2), \]  

(2)

where

\[ h_{X,LR}^\sigma(r_1, r_2) = \left| \sum_i^{N_\sigma} \psi_i(\sigma)(r_1) \psi_i(\sigma)(r_2) \right|^2 \frac{\rho_\sigma(r_1)}{\rho_\sigma(r_1)} \text{erf}(\omega r_{12}). \]  

(3)

The partitioning in Eq. 2 is then mirrored in the exchange energy, e.g. its long-range part reads

\[ E_{X,LR} = \sum_\sigma \frac{1}{2} \int \frac{\rho_\sigma(r_1) h_{X,LR}^\sigma(r_2)}{r_{12}} \text{d}r_1 \text{d}r_2. \]  

(4)

Regardless of the specific value of \( \omega \), modeling \( E_{X,LR} \) entirely by the orbital-dependent HF exchange constrains the asymptote of an exchange potential to the Coulombic decay, as in the exact theory,\(^{15}\)

\[ \hat{K}_X \psi_i(r) \xrightarrow{r \to \infty} -\frac{1}{r} \psi_i(r). \]  

(5)

where \( \psi_i(r) \) denotes an occupied orbital and \( \hat{K}_X \) is the exchange operator.

A system-specific \( \omega \) can be obtained non-empirically by means of optimal tuning.\(^{16}\) For instance, \( \omega \) can be adjusted so that the frontier (HOMO and LUMO) orbital energies, \( \epsilon_H \) and \( \epsilon_L \), match as closely as possible ionization potentials (IPs) and electron affinities (EAs), respectively.\(^{16,17}\) Another variant of this method, IP-tuning, constrains only the HOMO energy:\(^{18}\)

\[ J^2_H(\omega_H) = \min_\omega \left( E(N - 1) - E(N) + \epsilon_H \right)^2. \]  

(6)

All quantities on the right-hand side of Eq. 6 depend on \( \omega \). \( E(N) \) denotes the total energy of an \( N \)-electron system. Other choices of the objective function may also also be justified and are sometimes recommended.\(^{19}\) Eq. 6 imposes Koopmans’ theorem which is satisfied in the exact Kohn-Sham theory. Several authors demonstrated that tuning of an RS functional is a route towards reliable charge-transfer (CT) excitations,\(^{16,17,20}\) HOMO-LUMO (HL) gaps,\(^{5,16}\) photoelectron spectra,\(^{19}\) and hyperpolarizabilities.\(^{21}\) A survey of the
solutions of Eq. 6 for organic polymers\textsuperscript{18,22} reveals that $\omega_H$ dramatically depends on the system size and electronic structure.

The tuning procedure involves multiple evaluation of IPs, a step required to properly align single-particle spectra. In what follows, we will develop a model which enables a single-step computation of an approximation to the optimal RS parameter for a given system.

Theory

The physical picture behind the $-1/r$ asymptote of the exchange potential is an electron in the outer regions of a molecule interacting with its exchange hole, which is a compact charge distribution residing away from the reference electron, in the region of localized orbitals. (See Fig. 10 in Ref. 23 for an illustration of the exchange hole interacting with an electron beyond the last occupied shell.) Such an interaction is properly accounted for by the HF exchange hole. Consequently, the average separation of the outer-density electron from its exchange hole is an estimate of the interelectronic distance at which the long-range part of Eq. 1 should prevail. The average electron—exchange hole distance thus constitutes an upper bound on the value of $1/\omega$, which in turn is a measure of the distance at which the long-range part of Eq. 1 becomes dominant. To proceed further, we assume that this condition not only bounds $1/\omega$ from above, but completely defines the distance at which the transition between short- and long-range forms of the exchange hole occurs in an optimal RS functional.

We define the average squared electron—exchange hole distance as

$$
\langle d_{X,\sigma}^2 \rangle = \frac{\int \rho_\sigma(r)w_\sigma(r) d_{X,\sigma}^2(r) \, dr}{\int \rho_\sigma(r)w_\sigma(r) \, dr},
$$

(7)

where $d_{X,\sigma}$ is the vector pointing from the position of the reference electron to the charge center of the HF exchange hole, $\rho_\sigma$ is the $\sigma$-spin electron density, and $w_\sigma$ is a weight function. The weight function is required so that the density-tail region influences $\langle d_{X,\sigma}^2 \rangle$ more than
the bulk region. The choice of $w_{\sigma}$ is not unique. For instance, the formulas developed for switching between bulk and asymptotic exchange-correlation potentials in the context of asymptotic correction methods\textsuperscript{24,25} could be used. We decided to use a step function,

$$w_{\sigma}(r) = \begin{cases} 
1, & t_{\sigma}(r) \leq \mu, \\
0, & t_{\sigma}(r) > \mu,
\end{cases}$$  \hspace{1cm} (8)

which is based on a simplified variant of the electron localization function\textsuperscript{26} proposed by Schmider and Becke\textsuperscript{27},

$$t_{\sigma}(r) = \frac{\tau_{\text{UEG},\sigma}(r)}{\tau_{\sigma}(r)};$$  \hspace{1cm} (9)

$$\tau_{\text{UEG},\sigma} = \frac{3}{5} \left( 6\pi^2 \right)^{2/3} \rho_{\sigma}^{5/3};$$  \hspace{1cm} (10)

$$\tau_{\sigma} = \sum_{i} |\nabla \psi_{i\sigma}|^2.$$  \hspace{1cm} (11)

$t_{\sigma}$ is greater than 1 in regions where localized orbitals dominate, and tends to zero in the density tail.\textsuperscript{27} Thus, for a sufficiently small $\mu$, $w_{\sigma} = 1$ beyond any localized orbital shell, and $w_{\sigma} = 0$ elsewhere. Our preliminary results have shown that for most systems satisfactory results are obtained if the value of $\mu$ is so chosen that $N_{\sigma} = 1$ in the normalization integral,

$$N_{\sigma} = \int \rho_{\sigma}(r)w_{\sigma}(r;\mu)\,dr.$$  \hspace{1cm} (12)

The inequality $\mu \geq 0.07$ is enforced to prevent $\mu$ from attaining extremely small numerical values. To retain the size consistency of $\langle d_{X,\sigma}^2 \rangle$, $N_{\sigma}$ should be set to the number of noncovalently bound subsystems when $\langle d_{X,\sigma}^2 \rangle$ is computed for a noncovalent complex. The issue of the size consistency of system-specific RS functionals is discussed further in the text. The vector $d_{X,\sigma}$ in Eq. 7 points to the center of charge of the exact, orbital-dependent exchange hole, $h_{X}^\sigma$. It can be computed analytically at a cost no higher than any single-electron
integrand on a molecular grid,\textsuperscript{28}

\[
\mathbf{d}_{\mathbf{X},\sigma}(\mathbf{r}) = -\int h_{\mathbf{X}}^\sigma(\mathbf{r}, \mathbf{r}') \mathbf{r}' \, d\mathbf{r}' - \mathbf{r} = \left( \frac{1}{\rho_\sigma(\mathbf{r})} \sum_{ij} f_{ij\sigma} \psi_i\sigma(\mathbf{r}) \psi_j\sigma(\mathbf{r}) \right) - \mathbf{r},
\]

\text{(13)}

\[
f_{ij\sigma} = \int \mathbf{r}' \psi_i\sigma(\mathbf{r}') \psi_j\sigma(\mathbf{r}') \, d\mathbf{r}'.
\]

\text{(14)}

Using Eq. 7 we define the global-density-dependent (GDD) RS parameter,

\[
\omega_{\text{GDD}} = \frac{C}{\sqrt{\langle d_{\mathbf{X},\sigma}^2 \rangle}},
\]

\text{(15)}

where $C$ is a numerical constant depending on the underlying functional, the fraction of short-range HF exchange, and the weight function used in the definition of $\langle d_{\mathbf{X},\sigma}^2 \rangle$. It does not depend on a physical system. The parameter $C$ for any particular functional was determined via least-squares fitting of the model RS parameter, $\omega_{\text{GDD}}$, to the IP-tuned $\omega_H$. The fitting was performed within the set of molecules presented in Fig. 1, excluding only the outliers: the noble gases, Cl\textsubscript{2}, Cl\textsubscript{F}, and F\textsubscript{2}. No empirical or higher level theoretical data were involved in determining $C$. The spin label is dropped from the left-hand side of Eq. 15 because all systems studied in this work are closed-shell. In general case, however, $\omega_{\text{GDD}}$ would be spin-dependent.

By PBE(\omega) we denote an RS functional composed of PBE correlation,\textsuperscript{29} the short-range PBE exchange by Henderson et al.\textsuperscript{30}, and the long-range HF exchange. PBE(\omega) contains no short-range HF exchange. The definition of PBEh(\omega) is the same as PBE(\omega), except for 20\% of the short-range HF exchange. PBEh(0.20), also known as LRC-\omega PBEh, is the functional developed by Rohrdanz et al.\textsuperscript{31}. In our experience, PBE(0.40) performs almost identically to LC-\omega PBE proposed by Vydrov and Scuseria\textsuperscript{32}, although it differs in the derivation of the short-range exchange. The $C$ parameters used to compute the GDD-based RS parameters for PBE($\omega_{\text{GDD}}$) and PBEh($\omega_{\text{GDD}}$) are 0.90 and 0.75, respectively.
The value of $\omega_{\text{GDD}}[\rho]$ need not be evaluated fully self-consistently. We recommend the following protocol: (i) Perform a single point calculation using PBE(0.40) or PBEh(0.20). (ii) Use the converged density to compute $\omega_{\text{GDD}}[\rho]$ and provide a guess for the PBE($\omega_{\text{GDD}}$) or PBEh($\omega_{\text{GDD}}$) calculation. A single cycle of this procedure is sufficient for well-converged results. All numerical results presented in this work were generated in this way, except for those in Figs. 1 and 6, where densities from self-consistent IP-optimized calculations were used. All calculations employed def2-TZVPP basis unless specified otherwise.

Results and discussion

Refaely-Abramson et al. observed that the optimal RS parameter should decrease as the characteristic radius of a molecule grows. This trend is captured by $\omega_{\text{GDD}}$, which mimics the behavior of $\omega_{\text{H}}$ across the wide range of systems, from single atoms to multiple aromatic rings, see Fig. 1. The differences between $\omega_{\text{H}}$ and $\omega_{\text{GDD}}$ are small, except for halogen diatomics: F$_2$, Cl$_2$, and ClF. For these systems $\omega_{\text{GDD}}$ deviates substantially from $\omega_{\text{H}}$, but it is PBE($\omega_{\text{GDD}}$) which leads to a better agreement of $-\epsilon_{\text{H}}$ with the experimental IPs. The advantage of PBE($\omega_{\text{GDD}}$) stems from the fact that IPs are obtained directly from Koopmans’ theorem, without involving any energy differences of neutrals and ions. This is important, since for the halogen diatomics these energy differences are of poor quality. Overall, both PBE($\omega_{\text{H}}$) and PBE($\omega_{\text{GDD}}$) yield small mean-absolute-percentage deviations (MAPDs) of $\epsilon_{\text{H}}$ energies from experimental IPs, 2.5% and 2.1%, respectively. The signed errors of PBE($\omega_{\text{GDD}}$) and PBE($\omega_{\text{H}}$) are bracketed by two curves: one corresponding to the HF method, and the other to the PBEh(0.20) functional of Rohrdanz et al. Although the characteristic radius of a molecule is the decisive factor in most cases, there exists a special case of the alkali diatomics, Li$_2$ and Na$_2$, for which both GDD and IP-optimized RS parameters reach the smallest values in our set of molecules, even smaller than in a more extended naphthalene. Although the RS parameters of PBEh($\omega_{\text{GDD}}$) and PBE($\omega_{\text{GDD}}$) differ due to the varying amount of short-
range HF exchange, the performance of these methods with respect to Koopmans’ theorem is almost identical, see Figs. 2 and 1.

Figure 1: Comparison of IP-optimized and GDD-based RS parameters for the PBE(ω) functional (upper panel). Deviation of HOMO energies from experimental IPs\(^{36}\) (lower panel). \(\text{C}_{10}\text{H}_8\) denotes naphthalene.

Figure 2: Comparison of IP-optimized and GDD-based RS parameters for the PBEh(ω) functional (upper panel). Deviation of HOMO energies from experimental IPs\(^{36}\) (lower panel).

Having established that the GDD-based functionals yield HOMO energies close to the IPs, we now turn to the interpretation of LUMO energies. In practical applications, the total energy in an RS approximation is minimized with respect to orbitals and not with respect to the electron density, unlike traditional Kohn-Sham theory.\(^{37}\) Consequently, the
asymptote of the exchange operator depends on whether $\hat{K}_X$ acts on an occupied or virtual orbital. This is similar to the well-known property of the exchange operator in HF theory. Formally, such a functional is defined within generalized Kohn-Sham (GKS) theory.\textsuperscript{11} If we consider an electron in an $N$-electron system which is far from the nuclei and other electrons, at the distance $r \to \infty$, it feels the attraction from the nuclei and the repulsion from the remaining $N - 1$ electrons. In the RS DFT model, the electronic potential the electron feels is $(N - 1)/r$, where $N/r$ is contributed by the Coulomb operator and $-1/r$ by the exchange operator to compensate for the Coulombic self-interaction. If an electron is attached to the $N$-electron system, it feels the repulsion of the remaining $N$ electrons. Such a picture is properly captured in the GKS with the extra electron occupying LUMO: the Coulomb operator now contributes no self-interaction, therefore the exchange potential does not decay as $-1/r$ when acting on a virtual orbital. The interpretation of LUMO as an orbital of the attached electron is supported in GKS and is not supported by the traditional Kohn-Sham theory, where the exact exchange potential is multiplicative and always decays as $-1/r$.

It was proven by Yang et al.\textsuperscript{12} that for RS functionals within GKS framework, $\epsilon_L$ is the slope of $E(N)$ between a neutral and an anion,

$$ \left( \frac{\partial E}{\partial N} \right)^{(+)}_v = \epsilon_L. \quad (16) $$

Thus, if $E(N)$ is a piecewise-linear curve, $-\epsilon_L$ can serve as an approximation to the EA. A deviation from the straight line is called the many-electron self-interaction error\textsuperscript{13} (MSIE). Figs. 3 and 4 show that the MSIE of a conventional hybrid functional (PBE0\textsuperscript{38}) is reduced several times by introducing RS exchange in the fixed-$\omega$ PBEh(0.20) functional. Note that ClF has a positive EA, whereas NH$_3$ is bound only by the finite basis set. Further reduction of the error occurs when we use a system-dependent RS parameter instead of a fixed value. Therefore, both IP-tuned and GDD-based functionals are characterized by an
order-of-magnitude lower MSIE than the PBE0 functional.

To verify whether the GDD method provides physically meaningful LUMOs, the HL orbital gaps were calculated for nucleobases, a series of acenes, and perylene-3,4,9,10-tetracarboxylic dianhydride (PTCDA), see Fig. 5. The anions of DNA nucleobases and anthracene have finite lifetimes. Their negative EAs are estimated by computing $\epsilon_L$ in the finite basis set. The reference EAs were computed with the GW method (nucleobases) and coupled-cluster theory (anthracene). In all cases, EAs have only a minor effect on the magnitude of the fundamental gap. PBE($\omega_{\text{GDD}}$) and its short-range hybrid counterpart, PBEh($\omega_{\text{GDD}}$), yield HL gaps deviating from the reference theoretical fundamental gaps by merely 2.7% and 2.1% on average, respectively. The system-independent RS functionals, PBEh(0.20) and CAM-B3LYP, fail to achieve an uniform error distribution across all system sizes. The MAPDs are 4.4% and 19.5%, respectively.

![Figure 3: Deviation of $E(N)$ from straight line for ClF molecule.](image-url)
Figure 4: Deviation of $E(N)$ from straight line for NH$_3$ molecule.

Figure 5: Deviation of HL gaps from the theoretical fundamental gaps for DNA nucleobases$^{39,42,43}$ (uracil, thymine, guanine, cytosine, adenine), linear acenes$^{40,44}$ ($n = 2, \ldots, 6$), and PTCDA.$^{45}$ The RS parameters for PBE($\omega_{\text{GDD}}$) are given as annotations.

Both $\omega_{\text{H}}$ and $\omega_{\text{GDD}}$ decrease monotonically when computed for organic homopolymers of increasing length. In linear alkanes, $\omega_{\text{GDD}}$ closely follows $\omega_{\text{H}}$, see Fig. 6. In contrast to the alkanes, we have observed a discrepancy between $\omega_{\text{GDD}}$ and $\omega_{\text{H}}$ in polymers with unsaturated bonds. For linear acenes, a decrease of $\omega_{\text{GDD}}$ with the chain length is less steep than that of $\omega_{\text{H}}$. The discrepancy grows with the chain length; for hexacene $\omega_{\text{GDD}}$ is equal to 0.27 bohr$^{-1}$, whereas Körzdörfer et al.$^{18}$ report $\omega_{\text{H}}$ lower by 0.08 bohr$^{-1}$. In this system, the HL orbital gap calculated with PBE($\omega_{\text{H}}$) is 0.49 eV too low, whereas the PBE($\omega_{\text{GDD}}$) functional is in
excellent agreement with the reference CCSD(T) value,\textsuperscript{40,44} being only 0.04 eV too high. In chains of oligothiophenes containing up to 14 units, $\omega_{\text{GDD}}$ decreases less steeply than $\omega_{\text{H}}$ and saturates at a value nearly two times larger than $\omega_{\text{H}}$, see Fig. 7. For the shortest chains ($n = 2, \ldots, 5$), the IPs calculated with PBEh($\omega_{\text{GDD}}$) agree with the experimental data\textsuperscript{46} to within 0.2 eV. PBEh($\omega_{\text{H}}$) increasingly deviates from the experimental IPs as the chain lengthens. For a 14-unit oligothiophene, the IPs obtained with PBEh($\omega_{\text{GDD}}$) and PBEh(0.20) are closer to a rough estimate of the IP in an infinite oligothiophene chain\textsuperscript{46} than the IP obtained with PBEh($\omega_{\text{H}}$).

\begin{figure}[h]
\centering
\includegraphics[width=\textwidth]{figure6.png}
\caption{IP-optimized and GDD RS parameters for linear alkane chains, C$_{2n}$H$_{4n+2}$. Calculations employed def2-TZVP basis.\textsuperscript{33,34} The IP-optimized RS parameters and geometries were taken from Ref. 18.}
\end{figure}

Frontier orbital energies are related to the lowest CT excitation energy in adiabatic time-dependent DFT through Mulliken’s rule,\textsuperscript{1}

$$\Delta E_{\text{CT}} \rightarrow \text{IP} - \text{EA} - \frac{1}{R} = \epsilon_L - \epsilon_H - \frac{1}{R}, \quad (17)$$

where $R$ denotes a donor—acceptor separation. Because the GDD method was demonstrated to align orbital energies with IPs, we expect that it can accurately describe CT excitations. Indeed, data in Table 1 show that the GDD method recovers the gas-phase experimental CT excitation energies of aromatic donor (Ar)-tetracyanoethylene (TCNE) noncovalent
complexes at the accuracy similar to the GW-Bethe-Salpeter method and the optimally-tuned RS BNL functional. A system-independent RS functional, PBEh(0.20), is characterized by a five times larger MAD for these systems. Fig. 8 shows the distance dependence of CT excitations in \( \text{C}_2\text{H}_4\cdots\text{C}_2\text{F}_4 \) complex. The SAC-CI energies used as the reference were obtained with larger basis set and are considerably lower than the values previously reported by Tawada et al. All the RS functionals (LC-BLYP, CAM-B3LYP, PBEh(\( \omega \)GDD), and PBEh(\( \omega \)GDD)) as well as HF yield accurate energy differences relative to the values at 5.0 Å. However, the absolute values of excitation energies differ among the methods by several electronvolts, as seen in the lower panel of Fig. 8. The constant shift by which the methods differ can be attributed to different values of the HL gap in Eq. 17. Of all the tested methods, the GDD-based RS functionals are the closest to the benchmark curve.

Table 1: Energies and oscillator strengths (in parentheses) of CT transitions in gas phase Ar-TCNE complexes. All energies are given in electronvolts.

|        | PBE(\( \omega \)GDD) | PBEh(\( \omega \)GDD) | BNL \( ^{20} \) | GW \( ^{48} \)  | exp. \( ^{47} \) |
|--------|---------------------|-----------------------|---------------|---------------|----------------|
| toluene| 3.33(0.03)          | 3.23(0.03)            | 3.4           | 3.27          | 3.36(0.03)     |
| o-xylene| 3.36(0.03)        | 3.26(0.03)            | 3.0           | 2.89          | 3.15(0.05)     |
| benzene| 3.71(0.02)         | 3.60(0.02)            | 3.8           | 3.58          | 3.59(0.02)     |
| \( \text{C}_{10}\text{H}_{8} \) | 2.63(0.00)          | 2.52(0.00)            | 2.7           | 2.55          | 2.60(0.01)     |
| MAD    | 0.10                | 0.08                  | 0.12          | 0.10          |                |

Figure 7: RS parameters (upper panel) and IPs from Koopmans’ theorem (lower panel) for oligothiophene chains. Experimental IPs were taken from Ref. 46. The values of \( \omega_H \) were taken from Ref. 18.
One may expect that a method for which \(-\epsilon_H \approx \text{IP}\) should at the same time yield accurate energies of Rydberg transitions. To test this hypothesis, we gauged the performance of GDD-based functionals in a set of small molecules for which experimental energies of Rydberg transitions are available: butadiene, acetaldehyde, acetone, ethylene, isobutene, and formaldehyde. The respective MADs are presented in Fig. 9. The GDD computations employed 6-311(3+,3+)G** basis set.\(^{50}\) The transition assignments, experimental energies, and the results for LC-BLYP and CAM-B3LYP were taken from Ref. 51. We have found that, statistically, the GDD method does not lead to such a pronounced accuracy gain over the other methods as in the case of CT transitions. Even in the case of ethylene, where HOMO energies are within 0.1 eV of the experimental \(-\text{IP}\), the Rydberg transitions are systematically overestimated by about 0.3 eV by both PBE(\(\omega_{\text{GDD}}\)) and PBEh(\(\omega_{\text{GDD}}\)), see Table 2. We conclude that, in the case of Rydberg transitions, an approximate alignment of \(\epsilon_H\) with \(-\text{IP}\) prevents huge outlying absolute errors, but does not guarantee reduction of MADs.

Similarly to other approaches employing system-specific RS parameter, the GDD-based functionals are not size-consistent. The size consistency can be retained by an RS functional
Figure 9: MADs between TDDFT and experimental energies of Rydberg transitions. All Rydberg transitions compiled in the database by Caricato et al.\textsuperscript{51} were used.

Table 2: Rydberg excitations in ethylene. All energies are given in electronvolts. The results for CAM-B3LYP, LC-BLYP, and EOM-CCSD are taken from Ref. 51. The experimental vertical IP is taken from Ref. 36.

|       | PBE(ωGDD) | PBEh(ωGDD) | CAM-B3LYP | LC-BLYP | EOM-CCSD | exp. |
|-------|-----------|------------|-----------|---------|----------|------|
| 1B_{3u} | 7.61      | 7.49       | 6.89      | 7.41    | 7.28     | 7.11 |
| 1B_{1g}  | 8.20      | 8.10       | 7.48      | 8.04    | 7.93     | 7.80 |
| 1B_{2g}  | 8.37      | 8.23       | 7.54      | 8.21    | 7.96     | 7.90 |
| 2A_{g}   | 8.59      | 8.47       | 7.87      | 8.52    | 8.31     | 8.28 |
| 2B_{3u}  | 9.09      | 8.96       | 8.26      | 8.99    | 8.80     | 8.62 |
| 3B_{3u}  | 9.33      | 9.20       | 8.43      | 9.27    | 9.06     | 8.90 |
| 4B_{3u}  | 9.43      | 9.30       | 8.55      | 9.43    | 9.16     | 9.08 |
| 3B_{1g}  | 9.54      | 9.40       | 8.53      | 9.53    | 9.28     | 9.20 |
| 2B_{1u}  | 9.48      | 9.34       | 8.58      | 9.50    | 9.28     | 9.33 |
| 5B_{3u}  | 9.89      | 9.76       | 8.85      | 9.90    | 9.62     | 9.51 |
| MAD     | 0.38      | 0.25       | 0.48      | 0.31    | 0.11     |      |
| $-\epsilon_H$ | 10.78 | 10.63 | 10.68 |
only if $\omega$ is fixed for all systems, as in the PBEh(0.20) functional. However, fixed-$\omega$ functionals violate other constraints of the exact functional: Koopmans’ condition, the straight-line behavior of $E(N)$ between integers, and scaling to the high-density limit.\textsuperscript{52} In contrast, optimally-tuned as well as GDD-based functionals satisfy these conditions at least approximately. The choice between a fixed-$\omega$ and system-specific functional is dictated by the constraints which are important for a particular application. For example, fixed-$\omega$ functionals are preferred for computing enthalpies of formation,\textsuperscript{53} for which the size-consistency is crucial. On the other hand, there is numerical evidence that optimally-tuned functionals can be successfully applied to compute non-covalent binding energies,\textsuperscript{54} provided that the optimal RS parameter is determined for the dimer and the same value is also used for the monomers. Also CT excitations in non-covalent systems can be excellently described by the system-specific methods, despite their lack of size-consistency, see the results for Ar···TCNE complexes in Table 1 herein and in Ref. 20. In case of non-covalent interactions, it is even possible to eliminate the lack of size-consistency of optimally-tuned or GDD-based functionals altogether by applying different values of $\omega$ for each of the interacting species within SAPT(DFT) framework\textsuperscript{55} or subsystem approaches.\textsuperscript{56}

**Conclusions**

To summarize, we devised a model of the electron—exchange hole interaction in the outer density region that leads to straightforward predictions of the interelectron distance at which the nonlocal exchange should supersede the local DFA. The model spawns a class of approximations based on existing RS functionals. Two of such approximations, PBE($\omega_{\text{GDD}}$) and PBEh($\omega_{\text{GDD}}$), have been shown to align orbital energies with IPs and accurately describe CT excitations. The GDD method improves upon the existing approaches to the range separation in the following ways. (i) It is the first method capable of aligning single particle energies with IPs without the need for iterative calculation of ionized species. (ii) Unlike the
functionals with a fixed RS parameter, such as CAM-B3LYP and PBEh(0.20), its performance is not affected by the system size. (iii) $\omega_{\text{GDD}}$ depends on the global electron density and is constant in space, thus the computational cost is lower than in the local RS model of Krukau et al.\textsuperscript{52}. (iv) It offers superior quality of the frontier orbital energies in long unsaturated polymers such as oligothiophenes and acenes, for which the IP-tuning procedure predicts spuriously low values of the $\omega_{\text{H}}$ parameter.

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References

(1) Dreuw, A.; Weisman, J. L.; Head-Gordon, M. J. Chem. Phys. \textbf{2003}, \textit{119}, 2943–2946.

(2) Dreuw, A.; Head-Gordon, M. J. Am. Chem. Soc. \textbf{2004}, \textit{126}, 4007–4016.

(3) Tawada, Y.; Tsuneda, T.; Yanagisawa, S.; Yanai, T.; Hirao, K. J. Chem. Phys. \textbf{2004}, \textit{120}, 8425.

(4) Kamiya, M.; Tsuneda, T.; Hirao, K. J. Chem. Phys. \textbf{2002}, \textit{117}, 6010.

(5) Stein, T.; Eisenberg, H.; Kronik, L.; Baer, R. Phys. Rev. Lett. \textbf{2010}, \textit{105}, 266802.

(6) Gill, P. M.; Adamson, R. D. Chem. Phys. Lett. \textbf{1996}, \textit{261}, 105–110.

(7) Leininger, T.; Stoll, H.; Werner, H.-J.; Savin, A. Chem. Phys. Lett. \textbf{1997}, \textit{275}, 151–160.

(8) Iikura, H.; Tsuneda, T.; Yanai, T.; Hirao, K. J. Chem. Phys. \textbf{2001}, \textit{115}, 3540–3544.

(9) Becke, A. D. J. Chem. Phys. \textbf{1993}, \textit{98}, 1372–1377.
(10) Perdew, J. P.; Parr, R. G.; Levy, M.; Balduz Jr, J. L. Phys. Rev. Lett. 1982, 49, 1691–1694.

(11) Seidl, A.; Görling, A.; Vogl, P.; Majewski, J. A.; Levy, M. Phys. Rev. B 1996, 53, 3764–3774.

(12) Yang, W.; Cohen, A. J.; Mori-Sánchez, P. J. Chem. Phys. 2012, 136, 204111.

(13) Mori-Sánchez, P.; Cohen, A. J.; Yang, W. J. Chem. Phys. 2006, 125, 201102.

(14) Cohen, A. J.; Mori-Sánchez, P.; Yang, W. Chem. Rev. 2012, 112, 289–320.

(15) Baer, R.; Livshits, E.; Salzner, U. Ann. Rev. Phys. Chem. 2010, 61, 85–109.

(16) Kronik, L.; Stein, T.; Refaely-Abramson, S.; Baer, R. J. Chem. Theory Comput. 2012, 8, 1515–1531.

(17) Stein, T.; Kronik, L.; Baer, R. J. Chem. Phys. 2009, 131, 244119.

(18) Körzdörfer, T.; Sears, J. S.; Sutton, C.; Brédas, J.-L. J. Chem. Phys. 2011, 135, 204107.

(19) Refaely-Abramson, S.; Sharifzadeh, S.; Govind, N.; Autschbach, J.; Neaton, J. B.; Baer, R.; Kronik, L. Phys. Rev. Lett. 2012, 109, 226405.

(20) Stein, T.; Kronik, L.; Baer, R. J. Am. Chem. Soc. 2009, 131, 2818–2820.

(21) Sun, H.; Autschbach, J. ChemPhysChem 2013, 14, 2450–2461.

(22) Körzdörfer, T.; Parrish, R. M.; Sears, J. S.; Sherrill, C. D.; Brédas, J.-L. J. Chem. Phys. 2012, 137, 124305.

(23) Neumann, R.; Nobes, R.; Handy, N. Mol. Phys. 1996, 87, 1–36.

(24) Tozer, D. J.; Handy, N. C. J. Chem. Phys. 1998, 109, 10180–10189.

(25) Gruning, M.; Gritsenko, O. V.; van Gisbergen, S. J. A.; Baerends, E. J. J. Chem. Phys. 2001, 114, 652–660.
(26) Becke, A. D.; Edgecombe, K. E. J. Chem. Phys. 1990, 92, 5397–5403.

(27) Schmider, H.; Becke, A. J. Mol. Struct. THEOCHEM 2000, 527, 51–61.

(28) Becke, A.; Johnson, E. J. Chem. Phys. 2005, 122, 154104.

(29) Perdew, J.; Burke, K.; Ernzerhof, M. Phys. Rev. Lett. 1996, 77, 3865–3868.

(30) Henderson, T. M.; Janesko, B. G.; Scuseria, G. E. J. Chem. Phys. 2008, 128, 194105.

(31) Rohrdanz, M. A.; Martins, K. M.; Herbert, J. M. J. Chem. Phys. 2009, 130, 054112.

(32) Vydrov, O. A.; Scuseria, G. E. J. Chem. Phys. 2006, 125, 234109.

(33) Weigend, F.; Ahlrichs, R. Phys. Chem. Chem. Phys. 2005, 7, 3297–3305.

(34) Schuchardt, K. L.; Didier, B. T.; Elsethagen, T.; Sun, L.; Gurumoorthi, V.; Chase, J.; Li, J.; Windus, T. L. J. Chem. Inf. Model. 2007, 47, 1045–1052.

(35) Refaely-Abramson, S.; Baer, R.; Kronik, L. Phys. Rev. B 2011, 84, 075144.

(36) Lias, S.; Bartmess, J.; Liebman, J.; Holmes, J.; Levin, R.; Mallard, W. In NIST Chemistry WebBook, NIST Standard Reference Database Number 69; Linstrom, P., Mallard, W., Eds.; National Institute of Standards and Technology, 2013; Chapter Ion Energetics Data.

(37) Kohn, W.; Sham, L. J. Phys. Rev. 1965, 140, A1133–A1138.

(38) Adamo, C.; Barone, V. J. Chem. Phys. 1999, 110, 6158–6170.

(39) Faber, C.; Attaccalite, C.; Olevano, V.; Runge, E.; Blase, X. Phys. Rev. B 2011, 83, 115123.

(40) Hajgató, B.; Deleuze, M. S.; Tozer, D. J.; Proft, F. D. J. Chem. Phys. 2008, 129, 084308.
(41) Yanai, T.; Tew, D. P.; Handy, N. C. *Chem. Phys. Lett.* **2004**, *393*, 51–57.

(42) Bravaya, K. B.; Kostko, O.; Dolgikh, S.; Landau, A.; Ahmed, M.; Krylov, A. I. *J. Phys. Chem. A* **2010**, *114*, 12305–12317.

(43) Epifanovsky, E.; Kowalski, K.; Fan, P.-D.; Valiev, M.; Matsika, S.; Krylov, A. I. *J. Phys. Chem. A* **2008**, *112*, 9983–9992.

(44) Deleuze, M. S.; Claes, L.; Kryachko, E. S.; François, J.-P. *J. Chem. Phys.* **2003**, *119*, 3106–3119.

(45) Blase, X.; Attaccalite, C.; Olevano, V. *Phys. Rev. B* **2011**, *83*, 115103.

(46) Jones, D.; Guerra, M.; Favaretto, L.; Modelli, A.; Fabrizio, M.; Distefano, G. *J. Phys. Chem.* **1990**, *94*, 5761–5766.

(47) Hanazaki, I. *J. Phys. Chem.* **1972**, *76*, 1982–1989.

(48) Blase, X.; Attaccalite, C. *Appl. Phys. Lett.* **2011**, *99*, 171909.

(49) Nakatsuji, H.; Hirao, K. *J. Chem. Phys.* **1978**, *68*, 2053–2065.

(50) Caricato, M.; Trucks, G. W.; Frisch, M. J.; Wiberg, K. B. *J. Chem. Theory Comput.* **2010**, *7*, 456–466.

(51) Caricato, M.; Trucks, G. W.; Frisch, M. J.; Wiberg, K. B. *J. Chem. Theory Comput.* **2010**, *6*, 370–383.

(52) Krukau, A. V.; Scuseria, G. E.; Perdew, J. P.; Savin, A. *J. Chem. Phys.* **2008**, *129*, 124103.

(53) Karolewski, A.; Kronik, L.; Kummel, S. *J. Chem. Phys.* **2013**, *138*, 204115.

(54) Agrawal, P.; Tkatchenko, A.; Kronik, L. *J. Chem. Theory Comput.* **2013**, *9*, 3473–3478.
(55) Misquitta, A.; Podeszwa, R.; Jeziorski, B.; Szalewicz, K. J. Chem. Phys. 2005, 123, 214103.

(56) Rajchel, L.; Żuchowski, P. S.; Szczęśniak, M. M.; Chałasiński, G. Phys. Rev. Lett. 2010, 104, 163001.