Adiabatic connection from accurate wave-function calculations

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An extremely easy method for accurately calculating the adiabatic connection of density functional theory is presented, and its accuracy tested on both Hooke’s atom and the He atom. The method is easy because calculations are needed only for different values of parameters in the external potential, which can be achieved with almost any electronic structure code. Application of the method to accurate calculations on small systems should lead to benchmark adiabatic connection curves. © 2000 American Institute of Physics. [S0021-9606(00)30411-1]

I. INTRODUCTION

Density functional theory has become a popular computational method in quantum chemistry, because of its ability to handle large molecules accurately but relatively inexpensively. 1,2 This success is based on the availability of reliable accurate approximate functionals, and there is a constant need for still further accuracy. The goal of atomization energy errors being reliably less than 1 kcal/mol has not yet been achieved.

An important step forward in this search for accuracy came when Becke mixed a fraction of exact exchange with a generalized gradient approximation (GGA), and reduced errors by a factor of 2 or 3. 3 Such hybrid functionals, e.g., B3LYP, are now in common use, but their underlying justification comes from the adiabatic decomposition of density functional theory. Initially, the mixing parameters used were determined empirically. Later, it was shown that these parameters could be derived nonempirically. 4,5 and that a single universal mixing coefficient (25%) could be rationalized based on the performance of MP theory for molecules. 6 Most recently, new functionals have been proposed which use this adiabatic decomposition in much detail. 7

The adiabatic decomposition of an electronic system is very simple conceptually. Imagine multiplying the electron–electron repulsion by a coupling constant λ. Now imagine varying λ, while keeping the electron density ρ(r) fixed. This differs from traditional perturbative methods, e.g., Møller–Plesset, 8 because the external potential must be altered at each λ to keep the density fixed. At λ = 1, we have the physical, interacting electronic system. But as λ is reduced to zero, keeping the density fixed, the electrons become those of the noninteracting, Kohn–Sham system, and the potential morphs into the Kohn–Sham potential. All Kohn–Sham DFT calculations are actually performed on this noninteracting system, and the physical ground-state energy deduced from it. The adiabatic connection provides a continuous connection between the interacting system and its Kohn–Sham analog.

The only part of the total energy which must be approximated in such a Kohn–Sham calculation is the exchange–correlation energy, $E_{\text{XC}}(\rho)$, as a functional of the (spin) density. A further value of the coupling constant is that, through the Hellmann–Feynman theorem, this energy can be written as an integral over the purely potential contribution 9,10

$$E_{\text{XC}} = \int_0^1 d\lambda U_{\text{XC}}(\lambda), \quad U_{\text{XC}}(\lambda) = \langle \Psi^\lambda | V_{\text{ee}} | \Psi^\lambda \rangle - U,$$

(1)

where $V_{\text{ee}}$ is the Coulomb interaction between electrons, $\Psi^\lambda$ is the wave function at coupling constant λ, and $U$ is the Hartree electrostatic energy. This integral is the adiabatic connection formula. Hybrid functionals are based on the fact that this integrand, when applied to the exchange–correlation contribution to a bond dissociation energy, is often not well-approximated by GGA’s, due to their lack of static correlation. 11,12 This can be partially corrected for most molecules by mixing in a fraction of exact exchange. Construction of functionals based on this insight is often referred to as the adiabatic connection method (ACM).

Thus accurate approximation of this adiabatic connection curve is extremely important to further progress in construction of approximate functionals, and benchmark cases for small systems are always of interest and help in this endeavor. However, this is, in principle, a very demanding task. For each value of λ, one must solve the interacting electronic problem many times in order to find the external potential which reproduces the λ = 1 density. Almbladh and Pedroza 13 made early attempts, but not with the accuracy of modern calculations. In the last several years, as the importance of these curves has become apparent, several groups in different areas have performed adiabatic decomposition calculations. Hood et al. 14 have calculated the curve for bulk Si. Colonna and Savin 15 have used the slightly different Harris and Jones 16 decomposition for both the He and Be isoelectronic series. Joubert and Srivastava 17 have used Hylleras-type wave functions to calculate these curves for the He isoelectronic series.
The wave function, which we assume is a single Slater determinant of a function which has density \( \rho(r) \). In particular, \( \Psi_{\lambda=0}[\rho] \) is the noninteracting Kohn–Sham wave function, which we assume is a single Slater determinant. The noninteracting kinetic energy is

\[
T_{\lambda}[\rho] = \langle \psi_{\lambda=0} \mid \hat{T} \mid \psi_{\lambda=0} \rangle ,
\]

the Hartree electrostatic energy,

\[
U[\rho] = \frac{1}{2} \int d^3r \int d^3r' \frac{\rho(r) \rho(r')}{|r-r'|} ,
\]

and the exchange energy at coupling constant \( \lambda \) is

\[
E_{\lambda}[\rho] = \lambda \langle \psi_{\lambda=0} \mid \hat{V}_{xc} \mid \psi_{\lambda=0} \rangle - \lambda U[\rho] .
\]

In density functional theory, correlation energies are defined as differences between operators defined on correlated wave functions and the Kohn–Sham wave function. Thus the kinetic-correlation energy at \( \lambda \) is

\[
T_{xc}[\rho] = \langle \psi_{\lambda} \mid \hat{T} \mid \psi_{\lambda} \rangle - T_{\lambda}[\rho] ,
\]

while the potential-correlation energy at \( \lambda \) is

\[
U_{xc}[\rho] = \lambda \langle \psi_{\lambda} \mid \hat{V}_{xc} \mid \psi_{\lambda} \rangle - E_{\lambda}[\rho] ,
\]

so that the total correlation energy at \( \lambda \) is

\[
E_{xc}[\rho] = U_{xc}[\rho] + T_{xc}[\rho] .
\]

At \( \lambda = 1 \), the system is the true interacting quantum-mechanical system, and energies without superscripts refer to \( \lambda = 1 \), e.g., \( E_{xc} = E_{xc}^{\lambda=1} \), where the \( xc \) subscript implies the sum of the exchange and correlation energies.

All quantities at coupling constants different from one are simply related to their full coupling strength counterparts, but evaluated on a scaled density. The most important examples are the wave function

\[
\Psi_{\lambda}[\rho] = \Psi_{\lambda=0}[\rho_{\lambda=0}]
\]

and any exchange or correlation energies:

\[
E_{xc}^{\lambda}[\rho] = \lambda^2 E_{xc}[\rho_{\lambda=0}] ,
\]

where \( \rho_{\gamma}(r) = \gamma^2 \rho(\gamma r) \), and \( \Psi_{\gamma}(r_1 \cdots r_N) = \gamma^N \Psi(r_1 \cdots r_N) \). Thus knowledge of how a quantity varies as the density is scaled implies knowledge of its coupling constant dependence. These relations are trivial for energies evaluated on the Kohn–Sham wave function, since it is independent of the coupling constant.

\[
T_{\lambda}[\rho] = T_{\lambda=0}[\rho] \quad \text{or} \quad T_{\lambda}[\rho_{\gamma}] = \gamma^2 T_{\lambda}[\rho],
\]

the Hartree electrostatic energy

\[
U_{\gamma}[\rho] = \gamma U[\rho] ,
\]

and the exchange energy

\[
E_{\gamma}[\rho] = \gamma E_{\lambda}[\rho] \quad \text{or} \quad E_{\lambda}[\rho_{\gamma}] = \gamma E_{\lambda}[\rho] .
\]

Correlation is more sophisticated. Note, however, that knowledge of any quantity, \( E_{\lambda}[\rho_{\gamma}] / E_{\lambda}[\rho_{\gamma}] \) or \( U_{\lambda}[\rho_{\gamma}] \) as a function of \( \gamma \) between 1 and \( \infty \), i.e., scaling to the high density limit, is sufficient to determine the adiabatic connection for any of them, for \( \lambda \) between 0 and 1. The most well-known relation is to extract the kinetic-correlation piece from \( E_{\lambda}[\rho] \):

\[
T_{\lambda}[\rho] = -E_{\lambda}[\rho] + \left. \frac{dE_{\lambda}[\rho_{\gamma}]}{d\gamma} \right|_{\gamma=1} .
\]
To generalize this result to \( \rho_{\gamma} \), apply Eq. (14) to \( \rho_{\gamma} \), and make a change of variables in the derivative, to find

\[
T_C(\rho) = -E_C(\rho) + \gamma \frac{dE_C(\rho)}{d\gamma}.
\]  

(15)

This can be considered a first-order differential equation in \( \gamma \) for \( E_C(\rho_{\gamma}) \). Solution of this equation yields

\[
E_C(\rho_{\gamma}) = -\gamma \int_\gamma^\infty \frac{d\gamma'}{\gamma'} T_C(\rho_{\gamma'}).
\]  

(16)

where we have used the fact that \( E_C(\rho_{\gamma}) \) has vanished as \( \gamma \rightarrow 0 \). Via Eq. (10), these can be turned into coupling-constant relations [see Eq. (5) of Ref. 20]:

\[
T_S^\lambda = E_C^\lambda - \lambda \frac{dE_C^\lambda}{d\lambda},
\]

(17)

and

\[
E_C^\lambda = -\lambda \int_0^\lambda \frac{dN'}{\lambda^2} T_C^\lambda'.
\]

(18)

Similarly, we can extract \( U_C \) from \( E_C(\rho_{\gamma}) \), since \( U_C = E_C - T_C \). Thus Eq. (15) leads to

\[
U_C(\rho_{\gamma}) = 2E_C(\rho_{\gamma}) - \gamma \frac{dE_C(\rho_{\gamma})}{d\gamma},
\]

(19)

which, inverted, is

\[
E_C(\rho_{\gamma}) = \frac{\gamma}{2} \int_\gamma^\infty \frac{d\gamma'}{\gamma'} U_C(\rho_{\gamma'}).
\]

(20)

Combining Eqs. (19) and (16) then leads to simple relations between \( U_C \) and \( T_C \). Lastly, the coupling-constant relation that follows from Eq. (20), by applying Eq. (10) to \( E_C \) and \( U_C \), is

\[
E_C(\rho) = \int_0^1 \frac{d\lambda}{\lambda} E_C^\lambda(\rho),
\]

(21)

and \( U_{XC}(\lambda) = U_{XC}^\lambda/\lambda \) of Fig. 1 is just the integrand when this expression is applied to both exchange and correlation. Thus, the above reasoning may be considered a derivation of Eq. (1). We emphasize that all these relations follow from the theory of Ref. 20.

III. RESULTS

We present results for two systems, Hooke’s atom and the He atom. Hooke’s atom contains two electrons in a harmonic potential, interacting via a Coulomb repulsion:

\[
\hat{H} = -\frac{1}{2} \left( \nabla_1^2 + \nabla_2^2 \right) + \frac{1}{2|\mathbf{r}_1 - \mathbf{r}_2|} + \frac{1}{2} k(|\mathbf{r}_1^2 + \mathbf{r}_2^2|),
\]

(26)

and provides a valid test case for density functional theory, because of the Coulomb repulsion between electrons. It is also an easy system to perform calculations on, because the center-of-mass and relative coordinates separate, leaving only a one-dimension differential equation to solve numerically. Even this equation has an analytic solution for force constant \( k = 1/4 \), which is the system on which we will demonstrate most of our results.
To implement our method for the $k=1/4$ Hooke’s atom, we run many calculations at different values of $k>1/4$, up to $k=10$, using the method of Ref. 23. For each calculation, we scale the density so that it looks like the original $k=1/4$ density, using Eq. (22) to define the scale factor. Several such densities are shown in Fig. 2, which illustrates how close these densities are. Note that the largest error is for $\lambda = 0$, the noninteracting Gaussian density.

Figure 3 is a plot of the three quantities $E_C(\rho_{1/4})$, $T_C(\rho_{1/4})$, and $U_C(\rho_{1/4})$, all found using the PBE correlation functional. We choose these quantities to plot, as they remain close these densities are. Note that the largest error is for $\lambda = 0$, the noninteracting Gaussian density.

FIG. 2. Densities used for $\lambda = 1$ [least $\rho(0)$], 0.75, 0.5, 0.25, and 0 [largest $\rho(0)$].

FIG. 3. Correlation energies for the $k=1/4$ Hooke’s atom, calculated both with (solid lines) and without (dashed lines) the correction term, using the PBE correlation functional. The solid lines are indistinguishable from exact results, calculated by scaling the PBE correlation functional.

We also tested other possible prescriptions for choosing $\lambda$, such as from the square root of the ratio of noninteracting kinetic energy densities, as in Eq. (21). This gives values for $\lambda$ very close to those of Eq. (12), and leads to no measurable change in our estimate for $E_C(\rho_{1/4})$.

Figure 4 is the analog of Fig. 3, but now calculated using exact correlation energies and potentials. Based on the remarkable accuracy of Fig. 3, we claim these curves are essentially exact. In fact, the change due to the potential correction is much smaller for the exact case than for PBE, suggesting that the error made in Eq. (21) will also be smaller. The adiabatic curve of Fig. 1 was derived from these curves.

We also ran PBE calculations for lower densities, where the shape of the density can change significantly with $k$. We find that even as low as $k=10^{-4}$, there is a maximum error of only 1 millihartree in the corrected $E_C(\rho_{1/4})$ curve. Beyond this point, the reliability of our method might be questioned. However, defining an average $r_s$ value via Eq. (6a) of Ref. 25, at this point $\langle r_s \rangle = 19$, whereas for $k=1/4$, $r_s$ is 2.07. Thus, for common values of the density, the errors in our procedure are minute.

As a final test of our method, we return to the noninteracting kinetic energy. If the scaling were exact, the noninteracting kinetic energy would scale quadratically, as in Eq. (11). In Fig. 5, we plot $T_S[\rho_{s}] / \gamma^2$ exactly, approximating $T_S[\rho_{s}]$ by $T_S'(\rho')$, and including the correction of Eq. (25). Note that the maximum absolute error, at $\lambda = 0$, is only about $-1.6\%$, in the bare estimate. The correction makes the derivative with respect to $\lambda$ exact as $\lambda \to 1$, and overall reduces the error by about a factor of 5, to about $-0.3\%$.

Finally, we discuss our results for the two-electron ion series, simply to demonstrate that there is nothing special about Hooke’s atom which makes these techniques accurate. We may make bare estimates of the adiabatic curves directly from energy data already in the literature. Thus by calculating $\lambda$ by Eq. (22) using the data in Table I of Ref. 26, and assigning to each $\lambda$ the bare $U_C$ value from the table, we find the corresponding adiabatic connection curve integrates to $E_C = -41.5$ mH, as compared with an exact value of $-42.1$ mH for He. We can similarly use the $T_C$ data in Eq. (16) to calculate $T_C$.

similarity of densities, as shown in Fig. 2, leading to very small corrections.

We also tested other possible prescriptions for choosing $\lambda$, such as from the square root of the ratio of noninteracting kinetic energy densities, as in Eq. (21). This gives values for $\lambda$ very close to those of Eq. (12), and leads to no measurable change in our estimate for $E_C(\rho_{1/4})$.

FIG. 4. Correlation energies for the $k=1/4$ Hooke’s atom, calculated both with (solid lines) and without (dashed lines) the correction term.
get an estimate of $-41.2$ mH, so that the difference between these two results is a good indicator of the error in both of them. Again, we can calculate $T_S^1$, and find the largest error at $\lambda=0$, with a value of $2.916$, as compared to the exact value of $2.867$.

In Fig. 6, we plot PBE calculations of $E_C[\rho_{1/\alpha}]$ for the He atom, both exactly and using the adiabatic connection method described here, both with and without the correction term of Eq. (23). We use the highly accurate densities of Umrigar and Gonze. Unfortunately, not enough data points are available to make smooth plots for this system. Already at $Z=3, \lambda=0.62$, so that results for noninteger values of $Z$ (between $Z=2$ and $3$) are needed. However, enough data is present to see the clear correction the potential makes. We note several interesting features of this curve. First, the dependence on scaling is much less for He than for the $k=1/4$ Hooke’s atom. Second, the PBE curve contains a minimum, so Eq. (15) implies $E_C+T_C$ becomes positive for this scaled density. While it has never been rigorously proven, every known case of $E_C+T_C$ is negative. Thus this is probably a (very slight) limitation of PBE, which does not occur in PW91, and may be related to the difference at $r_g=0$ in Fig. 7 of Ref. 29. Third, the potential correction even picks up this feature, and still only has errors of a fraction of a millihartree.

![FIG. 5. Noninteracting kinetic energy as a function of coupling constant.](image1)

![FIG. 6. Correlation energy for the He atom, calculated both without (diamonds) and with (crosses) the correction term, using the PBE correlation functional. The solid line is the exact result, calculated by scaling the PBE correlation functional.](image2)

![FIG. 7. Correlation energy for the He atom, calculated both without (diamonds) and with (crosses) the correction term.](image3)

Figure 7 repeats Fig. 6, but for the exact case. Here the corrected curve is much straighter than for PBE, and has no minimum. The lines are drawn merely to aid the eye. Note that the value of $E^2_C[p]=\lim_{\rho \rightarrow \rho_g} E_C[\rho_g]$ drops about 1 millihartree, due to the difference in shape between the He atom and a pure exponential decay (bare estimate). We find $E^2_C= -47.9$ millihartree, in reasonable agreement with the value of Colonna and Savin (Ref. 15) ($-47.5$ from Table VI), of Joubert and Srivastava (Ref. 17) ($-47.2$ from $a_p$ in Table II), and of Engel and Dreizler (Ref. 30) ($-48.2$ from Table V). The value $-50.3$, reported in Ref. 31, was evaluated on the PBE self-consistent density.

**IV. CONCLUSIONS AND IMPLICATIONS**

We have shown how, with accurate ground-state results as a function of the external potential, an accurate adiabatic connection curve can be calculated for both Hooke’s atom at $k=1/4$ and the He atom. We see no reason why similar results could not be obtained for larger atoms, especially those for which the Kohn–Sham potential has already been calculated. There also exist methods for isolating the correlation potential. In the event that the exchange contribution cannot be easily isolated, the scheme can still be applied, but now to the combined exchange–correlation energy at each value of the external parameter. The correction is now evaluated using the exchange–correlation potential. On Hooke’s atom, this yields results almost as accurate as those described for the correlation energy alone. Important differences are in the regime $\lambda \rightarrow 0$, as here the exchange contribution to the correction blows up, since $\gamma=1/\lambda \rightarrow \infty$. This effect causes noticeable errors only for $\lambda<0.2$ in $E_C[\rho_{1/\alpha}]$, but not in $U_C$ or $T_C$. It also means that neither $U_C[\rho]$ or $T_C[\rho]$, as derived from the resulting $E_C[\rho_g]$ curve via Eqs. (19) and (15), is exact at $\lambda=1$. However, these differences are of the order of 0.1 millihartree.

Our method should work well for any system for which sufficiently accurate wave-function calculations are possible, and whose Kohn–Sham wave function is nondegenerate. We are currently investigating if this method can be used to calculate accurate adiabatic connection curves for other systems. Of particular interest would be a calculation for a large $Z$ four-electron ion, since the noninteracting hydrogenic
limit is degenerate in this case.\textsuperscript{35} Also of great interest are adiabatic connection curves for the binding energies of molecules,\textsuperscript{12} in which static correlation effects can be seen, and for which hybrid functionals can be justified.\textsuperscript{4,6} Due to the increased number of degrees of freedom, further investigation will be needed to determine the best procedure for approximating the scaled density in these cases.

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