

Fully numerical calculations on atoms with fractional occupations and range-separated exchange functionals

Susi Lehtola ^{*}*Department of Chemistry, University of Helsinki, P.O. Box 55 (A. I. Virtasen aukio 1), FI-00014 Helsinki, Finland*

(Received 17 October 2019; published 28 January 2020)

A recently developed finite-element approach for fully numerical atomic structure calculations [S. Lehtola, *Int. J. Quantum Chem.* **119**, e25945 (2019)] is extended to the description of atoms with spherically symmetric densities via fractionally occupied orbitals. Specialized versions of Hartree-Fock as well as local density and generalized gradient approximation density functionals are developed, allowing extremely rapid calculations at the basis-set limit on the ground and low-lying excited states, even for heavy atoms. The implementation of range separation based on the Yukawa or complementary error function (erfc) kernels is also described, allowing complete basis-set benchmarks of modern range-separated hybrid functionals with either integer or fractional occupation numbers. Finally, the computation of atomic effective potentials at the local density or generalized gradient approximation levels for the superposition of atomic potentials (SAP) approach [S. Lehtola, *J. Chem. Theory Comput.* **15**, 1593 (2019)] that has been shown to be a simple and efficient way to initialize electronic structure calculations is described. The present numerical approach is shown to afford beyond micro-Hartree accuracy with a small number of numerical basis functions, and to reproduce the literature results for the ground states of atoms and their cations for $1 \leq Z \leq 86$. Our results indicate that the literature values deviate by up to $10 \mu E_h$ from the complete basis-set limit. The numerical scheme for the erfc kernel is shown to work by comparison to results from large Gaussian basis-set calculations from the literature. Spin-restricted ground states are reported for Hartree-Fock and Hartree-Fock-Slater calculations with fractional occupations for $1 \leq Z \leq 118$.

DOI: [10.1103/PhysRevA.101.012516](https://doi.org/10.1103/PhysRevA.101.012516)

I. INTRODUCTION

Atoms are the simplest possible unit in chemistry, which is why electronic structure studies on atoms have a long and venerated history. Thanks to the high amount of symmetry that may be used to reduce the number of degrees of freedom in the atomic problem, fully numerical electronic structure approaches on atoms have been possible for a very long time [1]; for instance, a fully numerical configuration interaction calculation on the oxygen atom was reported by Hartree and co-workers over 80 years ago [2].

As the atomic Hamiltonian is spherically symmetric, the exact wave function should be rotationally invariant as well. Although the necessary symmetry requirements can straightforwardly be enforced in wave-function approaches, the application of density functional theory [3,4] (DFT) on atoms is surprisingly tricky. In the usual DFT approach, a single Slater determinant is employed, with all orbitals below the Fermi level being fully occupied. Nonrelativistically, all $2l + 1$ atomic orbitals sharing the principal quantum number n and angular quantum number l should be completely degenerate; however, this behavior is already broken by conventional DFT as well as Hartree-Fock (HF) on the first row. Different choices for the occupied orbitals on the $2p$ shell yield different final energies for, e.g., B and F, which may lead to several kcal/mol differences in the total energy—with a symmetrized density yielding yet another result [5]. One possibility to obtain comparable results is to employ a standard set of

electronic occupations [6], but such an approach does not yield the lowest possible energy.

Pursuing the lowest energy is not unproblematic, either. While HF is infamous for possessing variational solutions that break symmetry in systems with a high degree of symmetry [7], symmetry breaking is a problem in DFT as well [8]. In atoms, broken symmetries often arise for open shells, and the effect of nonspherical densities is known to be more pronounced with functionals at the generalized gradient approximation (GGA) level, and especially the meta-GGA (mGGA) level, than at the local density approximation (LDA) level [9–12]; even optimized effective potential exact-exchange calculations are subject to spurious energy splittings [13]. Inclusion of current density dependence leads to improvement of GGA and mGGA results [11,12], but the proper orbital degeneracy is still not fully restored.

Symmetry-breaking effects in atoms can already be seen at the simplest possible level of DFT, that is, the exchange-only LDA, which is also commonly known as Hartree-Fock-Slater (HFS) theory. For example, HFS calculations on the F atom reveal milli-Hartree decreases of the total energy upon addition of d as well as f functions, which is at variance to the generally accepted electronic configuration of fluorine as $1s^2 2s^2 2p^5$. Interestingly, this kind of symmetry breaking sometimes happens even in the case of closed-shell atoms; see, for instance, our recent finite-element reproduction [14] of calculations on atomic anions [15] where symmetry breaking was observed for H^- , Be, Li^- , and Na^- .

In addition to being degenerate due to symmetry (as often occurs in atoms), orbitals may also be degenerate by accident.

^{*}susi.lehtola@alumni.helsinki.fi

Since the aufbau rule implies populating the orbitals in increasing energy, it is tempting to divide the occupations evenly in the case of degeneracies. This paves the way for the use of fractional occupations, which in the case of atoms naturally yield a spherically symmetric density thanks to Unsöld's theorem [16]; the use of fractional occupations can be formally justified within the theory of ensemble representable densities [17–19].

Fractionally occupied orbitals should especially be used in the case where there is a negative gap between the highest occupied and lowest unoccupied orbital, no matter which way the orbitals are occupied; this happens when the highest occupied and lowest unoccupied orbital switch places during the orbital optimization. In this case, the total energy can be lowered by moving a fraction of an electron from the highest occupied orbital to the lowest unoccupied orbital, and at some point the two levels should cross.

Fractional occupations have been shown to yield better results for strongly correlated systems [20–24]. However, fractional occupations can only be justified at the Fermi level [25] and, more recently, it has been shown that energy minimization naturally leads to integer occupations below the Fermi level, and possible fractional occupations at the Fermi level for independent particle models such as HF and DFT [26].

While in some systems it is clear *a priori* from symmetry arguments or the orbital energies how many orbitals should be fractionally occupied, this is generally not the case. However, fractional occupations can be obtained as [27] the zero-temperature limit of finite-temperature DFT (FT-DFT) [28,29]. In a finite-temperature approach, the fractional orbital occupation numbers are determined by the orbital energies according to some smearing scheme that is typically controlled by a single parameter, i.e., an electronic temperature. Because of the simplicity and favorable computational scaling of FT-DFT, it has become a powerful tool for approximate modeling of systems exhibiting strong correlation; such approaches have been used to obtain promising results for a variety of systems [30–44].

Finite electronic temperatures may also be used to aid the convergence of self-consistent field calculations of molecules [45]; in the solid state, the use of fractional occupation numbers is often mandatory in order to attain convergence [46]. Although finite-temperature approaches are more attractive for DFT where all electrons experience the same potential, finite-temperature approaches can also be used in the context of HF calculations where they may offer good active spaces for post-HF calculations on strongly correlated systems [47].

Although several types of smearing schemes have been suggested, including Fermi-Dirac [28], Gaussian smearing [48], Methfessel-Paxton smearing [49], cold smearing [50], and others [51], they have been shown to yield similar results if the parameters are adjusted properly [47,52,53]; however, the behavior with respect to temperature needs to be carefully checked in each case to ensure convergence [54]. Note that the evaluation of forces in finite-temperature calculations requires the consideration of an additional entropic term that arises from the noninteger occupations and that depends on the smearing function [55,56].

Regardless of the used temperature, calculations with fractional occupations are more involved than those with integer occupations. Convergence acceleration techniques, such as direct inversion in the iterative subspace [57,58] (DIIS), become invalid when the orbital occupation pattern changes, even though the self-consistent field problem itself may become easier with fractional occupation numbers [45]. Determining the correct occupations is hard since the orbital occupations depend on the orbital energies, which in turn depend on the orbital occupations. The changes in the occupations may also cause changes in the shapes of the orbitals, meaning that the orbitals, their energies, and their occupations need to be solved self-consistently. Several approaches have been proposed for solving this problem both for zero [27,59–61] and finite electronic temperatures [51,62–66].

In systems with a high degree of symmetry such as atoms, the fractional occupations can be defined by symmetry block. Fractional occupations for atoms are typically defined in terms of atomic shells, over which the electrons are equally divided. For instance, the $1s^2 2s^2 2p^5$ configuration for F implies that the hole in the $2p$ shell be split in three, resulting in the minority-spin occupations $2p_x^{2/3} 2p_y^{2/3} 2p_z^{2/3}$; a spin-restricted variant would employ occupations of $2p_x^{5/6} 2p_y^{5/6} 2p_z^{5/6}$ in both spin channels. Indeed, this is the method of choice for fully numerical density functional calculations on atoms [1], and it has been used, e.g., in [67] for local density calculations on $1 \leq Z \leq 92$ at the ground-state electronic configuration from experiment, and in [68] for Perdew-Burke-Ernzerhof (PBE) [69,70] calculations on $1 \leq Z \leq 20$ and $31 \leq Z \leq 36$.

Atomic calculations with fractional occupation numbers are typically used to generate pseudopotentials [71,72], numerical atomic orbital basis functions [73,74], and Gaussian basis sets [75–77]. Spin-restricted spherically symmetric atoms may be used for setting up frozen-core calculations within all-electron approaches, and to determine approximate binding energies [78]. We have also recently shown that the radial potential from atomic calculations with fractional occupation numbers can be used to formulate efficient initial guesses for electronic structure calculations on polyatomic systems via the superposition of atomic potentials (SAP) approach [79].

In the typical case, electrons are divided evenly among the $2l + 1$ orbitals that are degenerate by symmetry. However, the fractional occupations can be generalized beyond integer occupations per shell, in case accidental degeneracy is also present. Early multiconfigurational HF calculations on atoms found that the $3d$ orbitals become occupied before the $4s$ orbitals in transition metals [80,81], which was solved by moving fractions of an electron between the shells. One example of this approach is the iron atom, where the $[\text{Ar}]3d^5 4s^1$ and $[\text{Ar}]3d^5_2 4s^1_0$ configurations both turn out to have a negative gap in the local density approximation [82], with the upper and lower indices denoting spin-up and spin-down electrons, respectively. With the Vosko-Wilk-Nusair (VWN) local density functional, the lowest-energy configuration is found to be $[\text{Ar}]3d^5_{1.3984} s^1_{0.602}$ [27].

A systematic, nonrelativistic study for spherical atoms in the range $1 \leq Z \leq 86$, has recently been presented by Kraissler *et al.* for the local density and PBE functionals based on

three local density functionals, employing 16 000 point grids and wave functions converged to $2 \mu E_h$ [83]. It was found in [83] that the ground state of most atoms does not involve fractional splitting of electrons between shells, indicating that a fully numerical program for modeling atoms with spherical densities would go a long way towards the final solution.

While several programs exist for either wave-function or density-functional-based fully numerical calculations on atoms [1], we are not aware of any publicly available software that supports hybrid functionals, except the recently published HELFEM program [14,84], which also includes a fully numerical approach for diatomic molecules that similarly supports hybrid functionals [85]. Most publicly available programs for fully numerical density functional calculations on atoms target the generation of projector-augmented wave (PAW) setups [86] or pseudopotentials [87]. Although Hartree-Fock pseudopotential generators have been available for some time [88,89], which allowed the use of non-self-consistent pseudopotentials for hybrid functionals [90], surprisingly, the self-consistent generation of pseudopotentials for hybrid functionals has only been described as of last year [91], explaining the scarcity of such programs.

Interestingly, the work of Yang *et al.* in [91] did not employ fractionally occupied Hartree-Fock calculations, but rather followed Slater's multiconfigurational approach, which is at odds with the density functional description used in the work, as the exact exchange and density functional parts experience different electron densities. In contrast, when fractional occupations are employed as in the present work, the exact exchange operator becomes independent of the magnetic quantum number m , as will be shown in Sec. II B, and both the density functional and exact exchange operators are evaluated with the same density matrix.

Although a general-use atomic program, such as the one in HELFEM, can be straightforwardly adapted to calculations on spherically symmetric densities by employing fractional occupation numbers in the construction of the density matrix, a more efficient approach is afforded by taking the assumption of the spherical symmetry of the density matrix deeper in the algorithms. As a result, some or even all of the angular integrals can be eliminated from the calculations, reducing the problem to a small number of dimensions; indeed, this is exactly what is done in the multiconfigurational HF approach Slater proposed 90 years ago [92].

In the present work, we describe the extension of the atomic program in HELFEM to the description of atoms with spherical symmetric density via fractional occupation numbers. Like the other programs in HELFEM, the spherically symmetric atomic program is interfaced to the LIBXC library of density functionals [93] and can be used with all supported density functionals therein. Specialized implementations for atomic calculations with fractional occupations are developed for local density approximation (LDA) and generalized gradient (GGA) functionals as well as HF exchange, yielding significant reductions in the dimensionality of the problem, whereas meta-GGA functionals can be used via an interface to the algorithms previously developed in [14].

Importantly, we also describe the implementation of Yukawa and complementary error function (erfc) range-separated exchange for atomic calculations in HELFEM with

either fractional or integer occupations, allowing complete basis-set benchmarks of recently developed exchange-correlation functionals such as the CAM-QTP family by Bartlett and co-workers [94–96], the N12-SX and revM11 functionals by Truhlar and co-workers [97,98], and the ω B97X-V and ω B97M-V functionals by Mardirossian and Head-Gordon (without the nonlocal correlation part) [99,100]. While the spherical harmonics decomposition for the Yukawa kernel is well known, the decomposition for the erfc kernel was only derived some time ago [101] and does not appear to have been implemented within a generally applicable, fully numerical approach for atoms. Results for H and He with relatively low-order B-spline basis sets have, however, been published almost simultaneously to our work [102]. Finally, we also describe the analytic calculation of the radial potentials necessary for the SAP orbital guess [79].

In Sec. II, we derive the equations for fractionally occupied HF and DFT at the LDA and GGA levels within the used finite-element approach. Then, in Sec. III, we present applications of the program to reproducing ground states for the neutral atoms and cations $1 \leq Z \leq 86$ and compare with [83]; we reproduce the long-range corrected density functional calculations on closed-shell atoms of [15] to show that the range-separation scheme works; and, finally, we report the nonrelativistic ground states of all atoms in the periodic table at HF and HFS levels of theory. The article concludes with a brief summary and discussion in Sec. IV.

II. METHOD

A basis set of the form

$$\chi_{nlm}(\mathbf{r}) = r^{-1} B_n(r) Y_l^m(\theta, \phi) \quad (1)$$

is adopted as in the integer-occupation program described in [14]. Here, $B_n(r)$ are the piecewise polynomial-shape functions of the finite-element method, which have been discussed extensively in [1] and [14] to which we refer for further details.

A. Range-separated exchange

As discussed in [1] and [14], the key to fully numerical electronic structure calculations on atoms is the Laplace expansion

$$\frac{1}{r_{12}} = \frac{4\pi}{r_{>}} \sum_{L=0}^{\infty} \frac{1}{2L+1} \left(\frac{r_{<}}{r_{>}}\right)^L \sum_{M=-L}^L Y_L^M(\Omega_1) [Y_L^M(\Omega_2)]^*, \quad (2)$$

which factorizes the two-electron integrals

$$(ij|kl) = \int \frac{\chi_i(\mathbf{r}) \chi_j^*(\mathbf{r}) \chi_k(\mathbf{r}') \chi_l^*(\mathbf{r}')}{|\mathbf{r} - \mathbf{r}'|} d^3r d^3r' \quad (3)$$

into a radial and an angular part.

In range-separated density functional theory [103,104], the Coulomb interaction is split into a short-range (sr) and a long-range (lr) part as

$$\begin{aligned} \frac{1}{r} &= \frac{\phi_{\text{sr}}(r)}{r} + \frac{1 - \phi_{\text{sr}}(r)}{r} = \frac{\phi_{\text{sr}}(r)}{r} + \frac{\phi_{\text{lr}}(r)}{r} \\ &= \frac{1 - \phi_{\text{lr}}(r)}{r} + \frac{\phi_{\text{lr}}(r)}{r}, \end{aligned} \quad (4)$$

where $\phi_{\text{sr}}(r) = 1 - \phi_{\text{r}}(r)$ is a splitting function. Typically, the short-range part is described using density functional theory and the long-range part with HF theory, but in practice many functionals employ more flexibility: for instance, the CAM-B3LYP functional [105] contains 19% short-range and 65% long-range exact exchange.

The evaluation of the range-separated exchange functionals is simple if one has access to the Green's function expansion of the range-separated kernel as

$$\frac{\phi_{\text{sr}}(r)}{r_{12}} = \sum_{L=0}^{\infty} \frac{4\pi}{2L+1} \mathcal{G}_L(r_{<}, r_{>}, \mu) \times \sum_{M=-L}^L [Y_L^M(\Omega_1)]^* Y_L^M(\Omega_2), \quad (5)$$

where $\mathcal{G}_L(r_{>}, r_{<}, \mu)$ is the Green's function, $r_{>}$ and $r_{<}$ are the greater and smaller of r_1 and r_2 , respectively, and μ is the range-separation parameter. The Green's function for the (unscreened) classical Coulomb interaction can be identified from Eq. (2) as

$$\mathcal{G}_L^{\text{Coulomb}}(r_{>}, r_{<}) = \frac{r_{<}^L}{r_{>}^{L+1}}. \quad (6)$$

The implementation of the integrals in HELFEM is based on the primitive integrals defined in [14] as

$$I_{ijkl}^L = \frac{4\pi}{2L+1} \int dr_1 dr_2 B_i(r_1) B_j(r_1) \times B_k(r_2) B_l(r_2) \mathcal{G}_L(r_{>}, r_{<}, \mu), \quad (7)$$

where $B_i(r)$ are the piecewise polynomial basis functions of Eq. (1).

1. Yukawa kernel

The Yukawa-screened [106] potential, $\phi^{\text{sr}}(r_{12}) = \exp(-\lambda r_{12})$, has a relatively well-known simple expansion,

$$\frac{e^{-\lambda|r-r'|}}{|r-r'|} = 4\pi\lambda \sum_{L=0}^{\infty} i_L(\lambda r_{<}) k_L(\lambda r_{>}) \times \sum_{M=-L}^L Y_L^M(\Omega_1) [Y_L^M(\Omega_2)]^*, \quad (8)$$

where i_L and k_L are regular and irregular modified Bessel functions that are regular at zero and infinity, respectively. Due to the separability, Yukawa-screened functionals are easy to handle in fully numerical approaches. Indeed, the Yukawa Green's function is employed in several recently developed linear scaling approaches for solving the HF or Kohn-Sham equations for bound orbitals in molecular systems via the Helmholtz kernel [107–111]. The Yukawa interaction is also straightforward to implement in calculations with Slater-type orbitals [112–114]. It turns out that Yukawa screening can be implemented with Gaussian-type orbitals in a rather straightforward manner [115], as analogous integrals also arise within r_{12} wave-function theory [116,117]. Such implementations are, however, rare at the moment, even though it has been claimed that Yukawa screening yields more accurate atomization and charge transfer excitation energies than erfc screening [118]. The Green's function for the Yukawa interaction can be

read from Eq. (8) as

$$\mathcal{G}_L^{\text{Yukawa}}(r_{>}, r_{<}, \lambda) = (2L+1)\lambda i_L(\lambda r_{<}) k_L(\lambda r_{>}). \quad (9)$$

As the Yukawa interaction factorizes in $r_{>}$ and $r_{<}$, it can be implemented in a similar fashion to the full Coulomb interaction, given by Eq. (6), along the lines of [14].

2. erfc kernel

Most range-separated functionals, however, are based on the complementary error function (erfc) kernel $\phi_{\text{sr}}(r) = \text{erfc}(\mu r)$. Such functionals are easy to implement in Gaussian-basis programs, requiring but simple modifications to the two-electron integrals [119,120], as well as plane-wave programs since the kernel has a simple Fourier transform which is strongly attenuated at large momentum. In contrast, the implementation of the erfc kernel is more complicated in real-space approaches. Fortunately, spherical harmonic expansions for the erfc Green's functions are available in the literature [101,121], but their form is more involved than that of the Yukawa function in Eq. (8). The main complication is that the Green's function does not factorize in $r_{<}$ and $r_{>}$, which means that two-dimensional quadrature is always required. In the approach of [101], new variables are introduced as $\Xi = \mu R$ and $\xi = \mu r$, and

$$\mathcal{G}_L(R, r; \mu) = \mu \Phi_L(\Xi, \xi), \quad (10)$$

where Φ_L is a scaled radial function given by

$$\Phi_n(\Xi, \xi) = F_n(\Xi, \xi) + \sum_{m=1}^n F_{n-m}(\Xi, \xi) \frac{\Xi^{2m} + \xi^{2m}}{(\Xi\xi)^m} H_n(\Xi, \xi), \quad (11)$$

$$F_n(\Xi, \xi) = \frac{2}{\sqrt{\pi}} \sum_{p=0}^n \left(-\frac{1}{4\Xi\xi}\right)^{p+1} \frac{(n+p)!}{p!(n-p)!} \times [(-1)^{n-p} e^{-(\xi+\Xi)^2} - e^{-(\xi-\Xi)^2}], \quad (12)$$

$$H_n(\Xi, \xi) = \frac{1}{2(\Xi\xi)^{n+1}} [(\Xi^{2n+1} + \xi^{2n+1}) \text{erfc}(\Xi + \xi) - (\Xi^{2n+1} - \xi^{2n+1}) \text{erfc}(\Xi - \xi)] \quad (13)$$

(note that the lower limit of the sum in Eq. (12) is incorrect in [101], where it reads $p = 1$ instead of $p = 0$). Equations (11) to (13) are numerically unstable in the short range, which is why when either $\xi < 0.4$, or $\Xi < 0.5$ and $0 < \xi < 2\Xi$ [101], the Green's function is evaluated with a Taylor expansion,

$$\Phi_n(\Xi, \xi) = \sum_k \frac{D_{n,k}(\Xi)}{\Xi^{n+1}} \xi^{n+2k}, \quad (14)$$

$$D_{n,0}(\Xi) = \text{erfc} \Xi + \frac{\exp(-\Xi^2)}{\sqrt{\pi}} (2\Xi^2)^{n+1} \sum_{m=1}^n \frac{(2\Xi^2)^{-m}}{(2n-2m-1)!}, \quad (15)$$

$$D_{n,k}(\Xi) = \frac{\exp(-\Xi^2)}{\sqrt{\pi}} (2\Xi^2)^{n+1} \frac{2n+1}{k!(2n+2k+1)} \times \sum_{m=1}^k \binom{m-k-1}{m-1} \frac{(2\Xi^2)^{k-m}}{(2n+2k-2m-1)!}, \quad k \geq 0. \quad (16)$$

Despite the lack of factorization of the erfc Green's function, its evaluation can be carried out analogously to the Coulomb and Yukawa kernels. The primitive integrals, given by Eq. (7), can be divided into two cases thanks to the finite support of the piecewise polynomial basis functions, as discussed in [14]. In an intraelement integral, both ij and kl are within the same element, whereas in an interelement integral, ij are in one element and kl are in another. In analogy to the scheme for Coulomb integrals discussed in [14], the interelement integrals are evaluated with N_{quad} quadrature points in both ij and kl , whereas the intraelement integrals employ N_{quad} points in ij , and the kl quadrature is split into N_{quad} intervals, all of which employ a fresh set of N_{quad} quadrature points.

B. Self-consistent field calculations with fractional occupations

It is well known that atomic orbitals can be written in the form

$$\psi_{nlm}(\mathbf{r}) = R_{nl}(r)Y_l^m(\hat{\mathbf{r}}). \quad (17)$$

Employing smeared occupations as

$$n(\mathbf{r}) = \sum_{n=1}^{\infty} \sum_{l=n-1}^{\infty} f_{nl} \sum_{m=-l}^l |\psi_{nlm}(\mathbf{r})|^2 \quad (18)$$

$$= \sum_{nl} f_{nl} \frac{(2l+1)R_{nl}^2(r)}{4\pi} = n(r), \quad (19)$$

where f_{nl} is the occupation number of all the $2l+1$ orbitals on the (n, l) shell, one immediately sees that the density matrix is diagonal in l and m ,

$$P_{\mu\nu}^{\sigma} = \delta_{l_{\mu}, l_{\nu}} \delta_{m_{\mu}, m_{\nu}} P_{\mu\nu}^{l_{\mu}; \sigma}, \quad (20)$$

and that the elements of the density matrix only depend on the value of l .

The spherical averaging yields huge simplifications for density functional calculations. Now, the density is only a function of the radial coordinate, and its gradient,

$$\nabla n = \partial_r n \hat{\mathbf{e}}_r, \quad (21)$$

only depends on the radial coordinate. Following the usual projective approach [14,122], the LDA and GGA matrix elements,

$$K_{\mu\nu}^{\text{xc}; \sigma} = \int \left\{ \frac{\delta f_{\text{xc}}}{\delta n_{\sigma}(\mathbf{r})} \phi_{\mu}(\mathbf{r}) \phi_{\nu}(\mathbf{r}) + \left[2 \frac{\delta f_{\text{xc}}}{\delta \gamma_{\sigma\sigma}(\mathbf{r})} \nabla \rho_{\sigma}(\mathbf{r}) + \frac{\delta f_{\text{xc}}}{\delta \gamma_{\sigma\sigma'}(\mathbf{r})} \nabla \rho_{\sigma'}(\mathbf{r}) \right] \cdot \nabla [\phi_{\mu}(\mathbf{r}) \phi_{\nu}(\mathbf{r})] \right\} d^3r, \quad (22)$$

become greatly simplified as only the radial terms are picked up, and as the same radial basis is used for all l, m ; see Eq. (1). Note, however, that meta-GGAs that depend on the kinetic-energy density cannot be handled in the same fashion, as the kinetic-energy density is not manifestly dependent only on the radial coordinate as discussed, e.g., in [123]. Like the exact exchange discussed below, the meta-GGA potential turns out to depend on the l channel. Meta-GGA functionals can be used in the present program via a fractional-occupation interface to the full atomic routines discussed in [14].

The Coulomb matrix arising from Eq. (2) trivially reduces to a single term as the spherically symmetric density only consists of a single $L=0, M=0$ component. Exact exchange—either with the full Coulomb form of Eq. (6) or the range-separated versions in Eqs. (9) and (10)—is a bit more complicated, as both the integrals and the density matrix carry a dependence on the orbital angular momenta in the well-known equation

$$K_{\mu\nu} = \sum_{\sigma\tau} (\mu\sigma | \nu\tau) P_{\sigma\tau}. \quad (23)$$

By employing the blocking of the density matrix given in Eq. (20), the exchange matrix can be written as

$$K_{\mu\nu}^{l_{\text{out}}} = \sum_{\sigma\tau} (\mu\sigma | \nu\tau) P_{\sigma\tau} = \sum_{L=|l_{\text{in}}-l_{\text{out}}|}^{l_{\text{in}}+l_{\text{out}}} I_{\mu\sigma\nu\tau}^L P_{\sigma\tau}^{l_{\text{in}}} \frac{1}{2l_{\text{out}}+1} \times \sum_{m_{\text{in}}=-l_{\text{in}}}^{l_{\text{in}}} \sum_{m_{\text{out}}=-l_{\text{out}}}^{l_{\text{out}}} G_{Ll_{\text{in}}, m_{\text{out}}}^{Mm_{\text{in}}, l_{\text{out}}} G_{Ll_{\text{in}}, m_{\text{out}}}^{Mm_{\text{in}}, l_{\text{out}}}, \quad (24)$$

where L is a coupled angular momentum with z projection, $M = m_{\text{out}} - m_{\text{in}}$. Rearranging the contractions, it is then seen that

$$K_{\mu\nu}^{l_{\text{out}}} = \sum_L I_{\mu\sigma\nu\tau}^L \left(\sum_{l_{\text{in}}} P_{\sigma\tau}^{l_{\text{in}}} \left[\frac{1}{2l_{\text{out}}+1} \times \sum_{m_{\text{in}}=-l_{\text{in}}}^{l_{\text{in}}} \sum_{m_{\text{out}}=-l_{\text{out}}}^{l_{\text{out}}} G_{Ll_{\text{in}}, m_{\text{out}}}^{Mm_{\text{in}}, l_{\text{out}}} G_{Ll_{\text{in}}, m_{\text{out}}}^{Mm_{\text{in}}, l_{\text{out}}} \right] \right), \quad (25)$$

where the evaluation is done from the inside-most bracket out.

C. Cusp condition

One way to diagnose atomic wave functions is the Kato-Steiner cusp condition [124,125],

$$C = -\frac{1}{2Z} \frac{d \ln n(r)}{dr} \Big|_{r=0} = -\frac{1}{2Z} \frac{n'(0)}{n(0)}, \quad (26)$$

which yields the value $C=1$ for the exact HF or density functional solution [126]. The electron density $n(r)$ at the nucleus was obtained in [14] via l'Hôpital's rule as

$$n(0) = P_{\mu\nu} \lim_{r \rightarrow 0} \frac{B_{\mu}(r)B_{\nu}(r)}{r^2} \quad (27)$$

$$= P_{\mu\nu} \lim_{r \rightarrow 0} \frac{\frac{d^2}{dr^2} B_{\mu}(r)B_{\nu}(r)}{\frac{d^2}{dr^2} r^2} \quad (28)$$

$$= P_{\mu\nu} B'_{\mu}(0)B'_{\nu}(0). \quad (29)$$

Its derivative at the nucleus also turns out to have a simple expression:

$$n'(0) = P_{\mu\nu} \lim_{r \rightarrow 0} \left[\frac{B'_{\mu}(r)B_{\nu}(r)}{r^2} + \frac{B_{\mu}(r)B'_{\nu}(r)}{r^2} - 2 \frac{B_{\mu}(r)B_{\nu}(r)}{r^3} \right] = P_{\mu\nu} \lim_{r \rightarrow 0} \left[\frac{\frac{d^3}{dr^3} B_{\mu}(r)B_{\nu}(r)}{\frac{d^2}{dr^2} r^2} - 2 \frac{\frac{d^3}{dr^3} B_{\mu}(r)B_{\nu}(r)}{\frac{d^3}{dr^3} r^3} \right] = P_{\mu\nu} B''_{\mu}(0)B'_{\nu}(0). \quad (30)$$

The value of the cusp is printed out at the end of all atomic calculations in HELFEM.

D. Effective radial potential for SAP

In the SAP approach discussed in [79], approximate orbitals for a molecule are obtained by diagonalizing an effective one-body Hamiltonian in an external potential obtained as a superposition of radial atomic potentials. Once the atomic ground state has been found with any supported method in HELFEM, including HF and hybrid and meta-GGA functionals, the radial effective potential for the SAP approach can be calculated based on any LDA or GGA functional. Extensions to the exact exchange, as in the optimized effective potential method [127], as well as generalized Kohn-Sham methods for the radial potentials from meta-GGA functionals are left for future work.

If the radial potential is self-consistent, i.e., the same functional is used for both the atomic orbitals and the potential, the SAP guess will reproduce the atomic orbitals exactly [79]. The atomic potential comprises Coulomb and exchange-

correlation contributions, the calculation of which is presented in the following.

1. Coulomb potential

Employing the Laplace expansion, given by Eq. (2), the Coulomb potential at a point \mathbf{r} for a spherically symmetric charge distribution is

$$V(\mathbf{r}) = \int_0^\infty \frac{1}{r_{>}} n(r') r^2 dr'. \quad (31)$$

Expressing the orbitals as in Eq. (17) yields potential matrix elements of the form

$$V_{ij}(r) = \int_0^\infty \frac{1}{r_{>}} B_i(r') B_j(r') dr' \quad (32)$$

$$= \frac{1}{r} \int_0^r B_i(r') B_j(r') dr' + \int_r^\infty \frac{1}{r'} B_i(r') B_j(r') dr', \quad (33)$$

and one gets three cases depending on whether or not r is inside the element where i and j reside. Let the element begin at r_b and end at r_e . Now,

$$V_{ij}(r) = \begin{cases} r^{-1} \int_{r_b}^{r_e} B_i(r') B_j(r') dr', & r > r_e, \\ r^{-1} \int_{r_b}^r B_i(r') B_j(r') dr' + \int_r^{r_e} r'^{-1} B_i(r') B_j(r') dr', & r_b < r < r_e, \\ \int_{r_b}^{r_e} r'^{-1} B_i(r') B_j(r') dr', & r < r_b. \end{cases} \quad (34)$$

Like the two-electron integrals discussed above, the in-element potential $r_b < r < r_e$ has to be evaluated by slices at every radial quadrature point $(r_0, r_1, \dots, r_{n-1})$,

$$\begin{aligned} \int_{r_b}^{r_{k-1}} B_i(r') B_j(r') dr' &= \int_{r_b}^{r_0} B_i(r') B_j(r') dr' \\ &+ \sum_{l=1}^{k-1} \int_{r_{l-1}}^{r_l} B_i(r') B_j(r') dr', \quad (35) \\ \int_r^{r_e} r'^{-1} B_i(r') B_j(r') dr' &= \int_{r_{n-1}}^{r_e} r'^{-1} B_i(r') B_j(r') dr' \\ &+ \sum_{l=k}^{n-1} \int_{r_{l-1}}^{r_l} r'^{-1} B_i(r') B_j(r') dr'. \end{aligned} \quad (36)$$

2. Exchange-correlation potential

The functional derivative satisfies

$$\delta E = E[n + \delta n] - E[n] = \int \frac{\delta E}{\delta n} \delta n d^3 r, \quad (37)$$

and so

$$\delta E = \int \left(\frac{\delta E}{\delta n} \delta n + \frac{\delta E}{\delta \nabla n} \delta \nabla n \right) d^3 r. \quad (38)$$

Integrating by parts, one gets

$$\int \frac{\delta E}{\delta \nabla n} \delta \nabla n d^3 r = \left[\frac{\delta E}{\delta \nabla n} \delta n d^3 r - \int \nabla \frac{\delta E}{\delta \nabla n} \delta n d^3 r \right] \quad (39)$$

$$= - \int \nabla \frac{\delta E}{\delta \nabla n} \delta n d^3 r, \quad (40)$$

from which one can identify

$$v(\mathbf{r}) = \frac{\delta E}{\delta n} - \nabla \cdot \frac{\delta E}{\delta \nabla n}. \quad (41)$$

Expressing the functional in terms of

$$\gamma^{\sigma\sigma'} = \nabla n^\sigma \cdot \nabla n^{\sigma'}, \quad (42)$$

one has

$$\frac{\delta}{\delta \nabla n} = \frac{\delta \gamma}{\delta \nabla n} \frac{\delta}{\delta \gamma} = 2 \nabla n \frac{\partial}{\partial \gamma}, \quad (43)$$

and so,

$$v(\mathbf{r}) = \frac{\delta E}{\delta n} - 2 \nabla \cdot \left(\frac{\delta E}{\delta \gamma} \nabla n \right) \quad (44)$$

or, for an open-shell system,

$$v^\sigma(\mathbf{r}) = \frac{\delta E}{\delta n^\sigma} - \nabla \cdot \left(2 \frac{\delta E}{\delta \gamma^{\sigma\sigma'}} \nabla n^\sigma + \frac{\delta E}{\delta \gamma^{\sigma\sigma'}} \nabla n^{\sigma'} \right), \quad (45)$$

where $\sigma \neq \sigma'$.

To guarantee accuracy, the gradient terms have to be evaluated analytically. Fortunately, there is only radial dependence, so the gradient

$$\nabla f = \left(\frac{\partial f}{\partial r}, 0, 0 \right) \quad (46)$$

can be replaced by a radial derivative, and the divergence with

$$\nabla \cdot \mathbf{A} = \left[\frac{1}{r^2} \frac{\partial}{\partial r} (r^2 A_r), 0, 0 \right] = \left(\frac{\partial}{\partial r} A_r + \frac{2A_r}{r}, 0, 0 \right). \quad (47)$$

TABLE I. Total energies of neutral atoms for calculations with the local spin density exchange functional and the Vosko-Wilk-Nusair correlation functional (VWN), and a Perdew-Burke-Ernzerhof exchange-correlation functional based on the VWN parametrization for the correlation energy of the homogeneous electron gas (PBE). The differences $\Delta E = E$ (present work) $- E$ [83] are calculated relative to the fully numerical values from [83] and are reported in micro-Hartree.

Z	E (VWN)	ΔE (VWN)	E (PBE)	ΔE (PBE)	Z	E (VWN)	ΔE (VWN)	E (PBE)	ΔE (PBE)
1	-0.478671	0	-0.499963	-0	44	-4439.044607	1	-4443.255631	-6
2	-2.834836	0	-2.893287	-1	45	-4683.360538	1	-4687.685035	-6
3	-7.343957	0	-7.462726	-0	46	-4935.368406	0	-4939.811859	-5
4	-14.447209	-0	-14.630525	-0	47	-5195.037351	1	-5199.600560	-6
5	-24.353614	0	-24.606283	-0	48	-5462.390982	1	-5467.070748	-5
6	-37.470031	0	-37.795116	-0	49	-5737.313809	-0	-5742.114862	-6
7	-54.136799	0	-54.537743	-0	50	-6019.972345	0	-6024.895821	-6
8	-74.527410	0	-75.003219	-1	51	-6310.419326	1	-6315.467009	-6
9	-99.114192	0	-99.668342	-1	52	-6608.650476	0	-6613.811930	-6
10	-128.233481	-0	-128.869661	-1	53	-6914.777857	1	-6920.056261	-6
11	-161.447625	-0	-162.176267	-1	54	-7228.856106	1	-7234.254478	-7
12	-199.139406	-0	-199.958820	-1	55	-7550.561866	0	-7556.083016	-7
13	-241.321156	-0	-242.236076	-1	56	-7880.111578	-0	-7885.752916	-7
14	-288.222945	0	-289.236535	-1	57	-8217.648931	91	-8223.406822	220
15	-340.005794	-0	-341.120757	-2	58	-8563.489711	880	-8569.364450	663
16	-396.743948	0	-397.951200	-1	59	-8917.715777	6966	-8923.707847	6539
17	-458.671463	-0	-459.976078	-2	60	-9280.405670	16518	-9286.515623	16052
18	-525.946195	0	-527.352025	-2	61	-9651.650420	13676	-9657.878495	13601
19	-598.206032	-0	-599.716752	-2	62	-10031.516930	6437	-10037.864135	6383
20	-675.742283	0	-677.355243	-2	63	-10420.023146	1	-10426.490411	-8
21	-758.685248	0	-760.397795	-3	64	-10817.148260	858	-10823.727509	285
22	-847.314902	-0	-849.129808	-3	65	-11223.108037	4860	-11229.800083	3513
23	-941.786662	-0	-943.704413	-2	66	-11637.977781	11030	-11644.783485	9215
24	-1042.218348	-0	-1044.239902	-3	67	-12061.832318	18521	-12068.752563	16445
25	-1148.644093	0	-1150.765417	-3	68	-12494.746152	26769	-12501.781831	24564
26	-1261.223291	6017	-1263.441835	4291	69	-12936.809752	19185	-12943.957962	20751
27	-1380.193787	716	-1382.508399	831	70	-13388.048594	1	-13395.317842	-9
28	-1505.672905	1	-1508.087914	-4	71	-13848.234767	1	-13855.623680	-9
29	-1637.793358	0	-1640.310279	-4	72	-14317.517965	2	-14325.032671	-9
30	-1776.573850	0	-1779.194575	-4	73	-14795.971453	1	-14803.612704	-9
31	-1921.851924	-0	-1924.582672	-3	74	-15283.610347	2	-15291.380462	-9
32	-2073.829860	0	-2076.672928	-4	75	-15780.381133	2	-15788.268506	-8
33	-2232.587154	0	-2235.545023	-4	76	-16286.434007	2	-16294.440422	-9
34	-2398.134930	0	-2401.196896	-4	77	-16801.850893	2	-16809.976281	-8
35	-2570.626651	-0	-2573.796934	-4	78	-17326.660985	3	-17334.912620	-10
36	-2750.147940	1	-2753.430126	-4	79	-17860.796573	2	-17869.175326	-10
37	-2936.342160	0	-2939.739646	-5	80	-18404.274220	1	-18412.777007	-10
38	-3129.453161	1	-3132.963153	-5	81	-18956.962102	1	-18965.593468	-10
39	-3329.525142	0	-3333.148098	-5	82	-19519.010773	2	-19527.771776	-10
40	-3536.771074	-0	-3540.515940	-5	83	-20090.453943	1	-20099.346370	-9
41	-3751.295618	0	-3755.160742	-6	84	-20671.273855	2	-20680.287630	-9
42	-3973.162595	-0	-3977.149787	-5	85	-21261.559507	2	-21270.697436	-10
43	-4202.348934	1	-4206.446961	-5	86	-21861.346869	3	-21870.611766	-9

Now,

$$\partial_r \left(\frac{\delta E}{\delta \gamma^{\sigma\sigma'}} \partial_r n^{\sigma'} \right) = \left(\partial_r \frac{\delta E}{\delta \gamma^{\sigma\sigma'}} \right) \cdot \partial_r n^{\sigma'} + \frac{\delta E}{\delta \gamma^{\sigma\sigma'}} \partial_r^2 n^{\sigma'}, \quad (48)$$

where

$$\partial_r \left(\frac{\delta E}{\delta \gamma^{\sigma\sigma'}} \right) = \frac{\partial n^\tau}{\partial r} \left(\frac{\partial}{\partial n^\tau} \frac{\delta E}{\delta \gamma^{\sigma\sigma'}} \right) + \frac{\partial \gamma^{\tau\tau'}}{\partial r} \left(\frac{\partial}{\partial \gamma^{\tau\tau'}} \frac{\delta E}{\delta \gamma^{\sigma\sigma'}} \right) = g^\tau \frac{\delta^2 E}{\delta n^\tau \delta \gamma^{\sigma\sigma'}} + (l^\tau g^{\tau'} + g^\tau l^{\tau'}) \frac{\delta^2 E}{\delta \gamma^{\tau\tau'} \delta \gamma^{\sigma\sigma'}}, \quad (49)$$

where we have defined $g^\tau = \partial_r n^\tau$ and $l^\tau = \partial_r^2 n^\tau$, and the extra $2A_r/r$ term from the divergence, given by Eq. (47), yielding

$$\frac{2}{r} \left(2 \frac{\delta E}{\delta \gamma^{\sigma\sigma}} g^\sigma + \frac{\delta E}{\delta \gamma^{\sigma\sigma'}} g^{\sigma'} \right). \quad (50)$$

TABLE II. Total energies of atomic cations for calculations with the local spin density exchange functional and the Vosko-Wilk-Nusair correlation functional (VWN), and a Perdew-Burke-Ernzerhof exchange-correlation functional based on the VWN parametrization for the correlation energy of the homogeneous electron gas (PBE). The differences $\Delta E = E$ (present work) $-E$ [83] are calculated relative to the fully numerical values from [83] and are reported in micro-Hartree.

Z	E (VWN)	ΔE (VWN)	E (PBE)	ΔE (PBE)	Z	E (VWN)	ΔE (VWN)	E (PBE)	ΔE (PBE)
1	0.000000	0	0.000000	0	44	-4438.767375	1	-4442.987586	-6
2	-1.941703	0	-1.993741	-1	45	-4683.055688	-0	-4687.391656	-6
3	-7.142818	-0	-7.257274	-0	46	-4935.023835	-0	-4939.477807	-6
4	-14.115512	-0	-14.299957	-0	47	-5194.755725	0	-5199.329663	-6
5	-24.038275	-0	-24.294185	-0	48	-5462.065582	1	-5466.758188	-6
6	-37.037413	0	-37.365761	-1	49	-5737.101461	1	-5741.909394	-6
7	-53.585407	-0	-53.989486	-1	50	-6019.697098	0	-6024.624904	-6
8	-74.016721	-0	-74.500466	-1	51	-6310.085269	0	-6315.135445	-6
9	-98.450427	0	-99.012379	-1	52	-6608.317272	0	-6613.492318	-6
10	-127.418114	-0	-128.061506	-1	53	-6914.378893	1	-6919.666895	-6
11	-161.250340	0	-161.979238	-2	54	-7228.394173	0	-7233.799075	-7
12	-198.855669	0	-199.679196	-1	55	-7550.416737	0	-7555.942187	-7
13	-241.100595	0	-242.016828	-2	56	-7879.920522	984	-7885.569221	813
14	-287.918773	-0	-288.932217	-1	57 ^a	-8217.455429	1545	-8223.221428	2463
15	-339.618451	-0	-340.732787	-2	58	-8563.294377	7617	-8569.175665	6993
16	-396.356540	0	-397.574834	-1	59	-8917.518819	20242	-8923.517325	19853
17	-458.184562	0	-459.496045	-2	60	-9280.230184	12512	-9286.344462	14304
18	-525.360439	-0	-526.770845	-2	61	-9651.482296	5894	-9657.715083	7570
19	-598.039506	-0	-599.553203	-2	62	-10031.348348	2185	-10037.700262	3563
20	-675.514035	-0	-677.132344	-3	63	-10419.820399	1	-10426.294380	-8
21	-758.442642	1812	-760.161910	1575	64	-10816.942313	782	-10823.528176	-8
22	-847.065015	2847	-848.886280	2949	65	-11222.899249	11493	-11229.597850	8739
23	-941.523838	0	-943.448454	-2	66	-11637.769436	20564	-11644.578605	20710
24	-1041.944126	0	-1043.972993	-3	67	-12061.641951	12969	-12068.562285	16381
25	-1148.368924	-0	-1150.502293	-3	68	-12494.571851	7644	-12501.608100	10480
26	-1260.927746	-0	-1263.157817	-3	69	-12936.633640	4018	-12943.786682	5267
27	-1379.896444	0	-1382.218969	-3	70	-13387.827982	1	-13395.103732	-10
28	-1505.370040	0	-1507.793428	-4	71	-13847.999554	2	-13855.396289	-9
29	-1637.485140	0	-1640.010907	-4	72	-14317.267886	1533	-14324.788056	2091
30	-1776.217890	0	-1778.850041	-3	73	-14795.705335	1	-14803.354716	-8
31	-1921.629140	-0	-1924.365546	-4	74	-15283.334783	2	-15291.113757	-9
32	-2073.533337	0	-2076.379628	-4	75	-15780.100605	2	-15788.005290	-8
33	-2232.220332	0	-2235.179888	-4	76	-16286.150952	2	-16294.166211	-8
34	-2397.770127	-0	-2400.845945	-4	77	-16801.535551	3	-16809.673022	-8
35	-2570.180737	-0	-2573.360358	-4	78	-17326.305178	1	-17334.567568	-10
36	-2749.623528	1	-2752.911995	-4	79	-17860.511437	0	-17868.901185	-10
37	-2936.183045	0	-2939.584557	-5	80	-18403.949940	1	-18412.465847	-10
38	-3129.240801	-0	-3132.756842	-5	81	-18956.753577	1	-18965.392384	-10
39	-3329.295616	-0	-3332.926478	-5	82	-19518.743995	2	-19527.509994	-10
40	-3536.524561	871	-3540.277393	131	83	-20090.133387	2	-20099.028863	-9
41	-3751.036938	0	-3754.910732	-5	84	-20670.954321	2	-20679.981706	-9
42	-3972.894262	-0	-3976.890719	-5	85	-21261.180969	2	-21270.328813	-8
43	-4202.074923	0	-4206.181883	-5	86	-21860.912366	3	-21870.184141	-9

^aAn incorrect value was reported in [83] for La⁺; see main text.

Thus, altogether, the radial exchange(-correlation) potential is given by

$$v_{xc}^{\sigma}(r) = \frac{\delta E}{\delta n^{\sigma}} - \nabla \cdot \left(2 \frac{\delta E}{\delta \gamma^{\sigma\sigma}} \nabla n^{\sigma} + \frac{\delta E}{\delta \gamma^{\sigma\sigma'}} \nabla n^{\sigma'} \right) = v_{\text{LDA}}^{\sigma}(r) - \frac{2}{r} \left[2 \frac{\delta E}{\delta \gamma^{\sigma\sigma}} g^{\sigma} + \frac{\delta E}{\delta \gamma^{\sigma\sigma'}} g^{\sigma'} \right] \quad (51)$$

$$- 2 \left[g^{\tau} \frac{\delta^2 E}{\delta n^{\tau} \delta \gamma^{\sigma\sigma}} g^{\sigma} + (l^{\tau} g^{\tau'} + g^{\tau} l^{\tau'}) \frac{\delta^2 E}{\delta \gamma^{\tau\tau'} \delta \gamma^{\sigma\sigma}} g^{\sigma} + \frac{\delta E}{\delta \gamma^{\sigma\sigma}} l^{\sigma} \right] \\ - \left[g^{\tau} \frac{\delta^2 E}{\delta n^{\tau} \delta \gamma^{\sigma\sigma'}} g^{\sigma'} + (l^{\tau} g^{\tau'} + g^{\tau} l^{\tau'}) \frac{\delta^2 E}{\delta \gamma^{\tau\tau'} \delta \gamma^{\sigma\sigma'}} g^{\sigma'} + \frac{\delta E}{\delta \gamma^{\sigma\sigma'}} l^{\sigma'} \right], \quad (52)$$

where the various derivatives of the exchange-correlation functional are available in LIBXC [93].

III. RESULTS

To demonstrate the routines, we reproduce the literature values for the ground states of the neutral and cationic atoms, $1 \leq Z \leq 86$, with the VWN functional, accurate values for which were obtained in [83] with 10 000 radial grid points. A Perdew-Burke-Ernzerhof (PBE) [69,70] functional variant described in [83] is also considered, where the correlation energy in the homogeneous electron gas limit is given by the VWN parametrization instead of the Perdew-Wang [128] parametrization of the original PBE correlation functional.

We found that by using a radial basis consisting of ten 15-node elements and a practical infinity $r_\infty = 40a_0$, the energy has converged beyond micro-Hartree accuracy, even though this basis contains just 139 radial basis functions—almost two orders of magnitude fewer degrees of freedom than used in [83]. These results, with differences to the reference data from [83], are shown in Table I for neutral atoms and Table II for their cations.

The agreement, for the most part, is excellent: large positive differences that indicate that the value of [83] is lower are seen for the species for which the calculations in [83] transferred fractional charge across shells. Otherwise, the differences become noticeable for heavier atoms, nearing $-10 \mu E_h$ when $Z \rightarrow 86$, indicating that the data in [83] are not fully converged to the basis-set limit. In our VWN calculation on the La^+ cation, it was discovered that the energy for the $4f_0^{0.945}5d_0^{0.575}6s_0^{0.480}$ state reported in [83] was incorrect; the correct energy is -8217.456974 [129].

Next, we demonstrate that the erfc range-separation scheme works by reproducing the literature values for the total energies of the spherically symmetric atoms on the first two rows using the long-range corrected BLYP functional [15,130,131]. In [14], we investigated the accuracy of the aug-pc- ∞ Gaussian basis set that was used in [15]. The study was restricted to nS states to avoid symmetry-breaking effects, which were still observed for H^- , He , Li^- , and Na^- , as was discussed in Sec. I. Reproducing symmetry-preserving data with ERKALE [132], we found that the truncation error of the aug-pc- ∞ basis set is less than $1 \mu E_h$ for light atoms and tens of μE_h for heavier atoms in Hartree-Fock and BHLYP [133] calculations.

Because the screening is evaluated analytically in Gaussian-basis calculations [119] and the accuracy of the aug-pc- ∞ basis set has been established [14], the values reported in [15] offer an ideal reference for the present work. The comparison of results obtained in the present work with Eqs. (11) to (16) and a numerical basis set with five 15-node radial elements and a practical infinity $r_\infty = 40a_0$ is shown in Table III, demonstrating excellent agreement between the fully numerical and Gaussian-basis calculations. The values are in full agreement after rounding to the same accuracy for the light atoms, while the fully numerical values are slightly below the Gaussian-basis values for the heavier atoms, as expected based on the basis-set truncation errors observed in [14].

Finally, the spin-restricted ground states for all atoms in the periodic table at HF and HFS levels of theory are

TABLE III. Comparison of the total energies of spherically symmetric atoms with the LC-BLYP functional with the range separation constant $\omega = 0.3$ reproduced with finite-element calculations (present work) and a Gaussian basis-set calculation [15].

Atom	Finite element	Gaussian basis
H^-	-0.519949	-0.51995
He	-2.866811	-2.86681
Li^-	-7.435511	-7.43551
Be	-14.584723	-14.58472
N	-54.482223	-54.48222
F^-	-99.766050	-99.76604
Ne	-128.816627	-128.81661
Na^-	-162.136564	-162.13655
Mg	-199.907036	-199.90702
P	-341.069932	-341.06992
Cl^-	-460.080588	-460.08057
Ar	-527.321257	-527.32124

shown in Tables IV and V, respectively; these calculations also used ten 15-node radial elements. The data reveal that in some cases, a lower-lying configuration has been seen in the brute-force search, but that it failed to converge. In the HF calculations, the $6s^24f^25d^1$ state of Pr converges without problems; however, the $6s^24f^3$ configuration has a lower energy, but its wave function failed to converge. Similar issues are observed in the HFS calculations for Cf, Es, and Fm, where the $5f^{n-1}7s^1$ state converges without problems, but a lower energy is observed for a $5f^n$ configuration, the wave function of which fails to converge.

The HF results can be compared to the high-accuracy data for multiconfigurational HF of Saito [134]. Because the present calculations are spin restricted with fractional occupations, the energies are higher than those reported in [134]. However, the agreement for the noble gases is perfect, underlining the high accuracy of the computational approach used in the present work, which was outlined in [14], even though only 139 radial basis functions were employed.

IV. SUMMARY AND DISCUSSION

We have described efficient implementations of range-separated functionals as well as fractional occupations for atomic electronic structure calculations with HELFEM, and demonstrated that beyond micro-Hartree accuracy can be achieved with just 139 numerical radial basis functions. We have tested the program by reproducing local density approximation (LDA) and generalized gradient approximation (GGA) total energies for $1 \leq Z \leq 86$ at the basis-set limit, and shown that the literature values deviate from the complete basis-set limit by up to $10 \mu E_h$. The approaches developed in the present work could be straightforwardly extended to fractional occupations per shell in future work, requiring the addition of a logic to formulate the fractional occupation numbers.

The capabilities added to HELFEM in the present work allow for self-consistent benchmarking of density functionals at the basis-set limit, which is useful for development

TABLE IV. Nonrelativistic spin-restricted spherical HF configurations for all elements in the periodic table. Entries in italics indicate that a lower-lying configuration was identified but it failed to converge.

H	$1s^1$	-0.357710	Nb	$[\text{Kr}]5s^24d^3$	-3753.033166	Tl	$[\text{Xe}]4f^{14}5d^{10}6s^26p^1$	-18961.740923
He	$1s^2$	-2.861680	Mo	$[\text{Kr}]4d^6$	-3974.815043	Pb	$[\text{Xe}]4f^{14}5d^{10}6s^26p^2$	-19523.831389
Li	$[\text{He}]2s^1$	-7.378133	Tc	$[\text{Kr}]4d^7$	-4204.141902	Bi	$[\text{Xe}]4f^{14}5d^{10}6s^26p^3$	-20095.328128
Be	$[\text{He}]2s^2$	-14.573023	Ru	$[\text{Kr}]4d^8$	-4441.038680	Po	$[\text{Xe}]4f^{14}5d^{10}6s^26p^4$	-20676.283998
B	$[\text{He}]2s^22p^1$	-24.384693	Rh	$[\text{Kr}]4d^9$	-4685.600291	At	$[\text{Xe}]4f^{14}5d^{10}6s^26p^5$	-21266.749081
C	$[\text{He}]2s^22p^2$	-37.344157	Pd	$[\text{Kr}]4d^{10}$	-4937.921024	Rn	$[\text{Xe}]4f^{14}5d^{10}6s^26p^6$	-21866.772241
N	$[\text{He}]2s^22p^3$	-53.852155	Ag	$[\text{Kr}]4d^{10}5s^1$	-5197.639939	Fr	$[\text{Rn}]7s^1$	-22475.826522
O	$[\text{He}]2s^22p^4$	-74.297532	Cd	$[\text{Kr}]4d^{10}5s^2$	-5465.133143	Ra	$[\text{Rn}]7s^2$	-23094.303666
F	$[\text{He}]2s^22p^5$	-99.067145	In	$[\text{Kr}]4d^{10}5s^25p^1$	-5740.082317	Ac	$[\text{Rn}]7s^26d^1$	-23722.073196
Ne	$[\text{He}]2s^22p^6$	-128.547098	Sn	$[\text{Kr}]4d^{10}5s^25p^2$	-6022.746221	Th	$[\text{Rn}]7s^26d^2$	-24359.362900
Na	$[\text{Ne}]3s^1$	-161.808533	Sb	$[\text{Kr}]4d^{10}5s^25p^3$	-6313.211503	Pa	$[\text{Rn}]7s^25f^3$	-25006.406325
Mg	$[\text{Ne}]3s^2$	-199.614636	Te	$[\text{Kr}]4d^{10}5s^25p^4$	-6611.551696	U	$[\text{Rn}]7s^25f^4$	-25663.398242
Al	$[\text{Ne}]3s^23p^1$	-241.782323	I	$[\text{Kr}]4d^{10}5s^25p^5$	-6917.837495	Np	$[\text{Rn}]7s^25f^5$	-26330.321976
Si	$[\text{Ne}]3s^23p^2$	-288.637472	Xe	$[\text{Kr}]4d^{10}5s^25p^6$	-7232.138364	Pu	$[\text{Rn}]5f^77s^1$	-27007.271797
P	$[\text{Ne}]3s^23p^3$	-340.381142	Cs	$[\text{Xe}]6s^1$	-7553.899845	Am	$[\text{Rn}]5f^9$	-27694.356363
S	$[\text{Ne}]3s^23p^4$	-397.202080	Ba	$[\text{Xe}]6s^2$	-7883.543827	Cm	$[\text{Rn}]5f^{10}$	-28391.573019
Cl	$[\text{Ne}]3s^23p^5$	-459.286063	La	$[\text{Xe}]6s^25d^1$	-8220.935691	Bk	$[\text{Rn}]5f^{11}$	-29098.977559
Ar	$[\text{Ne}]3s^23p^6$	-526.817513	Ce	$[\text{Xe}]6s^25d^2$	-8566.342481	Cf	$[\text{Rn}]5f^{12}$	-29816.624759
K	$[\text{Ar}]4s^1$	-599.124244	Pr	$[\text{Xe}]6s^24f^25d^1$	-8920.094872	Es	$[\text{Rn}]5f^{13}$	-30544.570349
Ca	$[\text{Ar}]4s^2$	-676.758186	Nd	$[\text{Xe}]6s^24f^4$	-9282.434373	Fm	$[\text{Rn}]5f^{14}$	-31282.870930
Sc	$[\text{Ar}]4s^24p^1$	-759.556762	Pm	$[\text{Xe}]6s^24f^5$	-9653.359914	Md	$[\text{Rn}]5f^{14}7s^1$	-32031.135295
Ti	$[\text{Ar}]4s^23d^2$	-847.933865	Sm	$[\text{Xe}]6s^24f^6$	-10032.949725	No	$[\text{Rn}]5f^{14}7s^2$	-32789.512140
V	$[\text{Ar}]4s^23d^3$	-942.147322	Eu	$[\text{Xe}]4f^76s^2$	-10421.286649	Lr	$[\text{Rn}]5f^{14}7s^26d^1$	-33557.812903
Cr	$[\text{Ar}]4s^23d^4$	-1042.342957	Gd	$[\text{Xe}]6s^14f^9$	-10818.487373	Rf	$[\text{Rn}]5f^{14}7s^26d^2$	-34336.316816
Mn	$[\text{Ar}]3d^7$	-1148.803487	Tb	$[\text{Xe}]4f^{11}$	-11224.646666	Db	$[\text{Rn}]5f^{14}7s^26d^3$	-35125.088022
Fe	$[\text{Ar}]3d^8$	-1261.579698	Dy	$[\text{Xe}]4f^{12}$	-11639.819030	Sg	$[\text{Rn}]5f^{14}6d^6$	-35924.293864
Co	$[\text{Ar}]3d^9$	-1380.817569	Ho	$[\text{Xe}]4f^{13}$	-12064.074984	Bh	$[\text{Rn}]5f^{14}6d^7$	-36733.871607
Ni	$[\text{Ar}]3d^{10}$	-1506.669759	Er	$[\text{Xe}]4f^{14}$	-12497.495312	Hs	$[\text{Rn}]5f^{14}6d^8$	-37553.863992
Cu	$[\text{Ar}]3d^{10}4s^1$	-1638.899667	Tm	$[\text{Xe}]4f^{14}6s^1$	-12939.976389	Mt	$[\text{Rn}]5f^{14}6d^9$	-38384.313942
Zn	$[\text{Ar}]3d^{10}4s^2$	-1777.848116	Yb	$[\text{Xe}]4f^{14}6s^2$	-13391.456193	Ds	$[\text{Rn}]5f^{14}6d^{10}$	-39225.264332
Ga	$[\text{Ar}]3d^{10}4s^24p^1$	-1923.166449	Lu	$[\text{Xe}]4f^{14}6s^26p^1$	-13851.687533	Rg	$[\text{Rn}]5f^{14}6d^{10}7s^1$	-40076.301440
Ge	$[\text{Ar}]3d^{10}4s^24p^2$	-2075.150884	Hf	$[\text{Xe}]4f^{14}6s^25d^2$	-14320.929628	Cn	$[\text{Rn}]5f^{14}6d^{10}7s^2$	-40937.797856
As	$[\text{Ar}]3d^{10}4s^24p^3$	-2233.924574	Ta	$[\text{Xe}]4f^{14}6s^25d^3$	-14799.321729	Nh	$[\text{Rn}]5f^{14}6d^{10}7s^27p^1$	-41809.456590
Se	$[\text{Ar}]3d^{10}4s^24p^4$	-2399.595885	W	$[\text{Xe}]4f^{14}5d^6$	-15286.959470	Fl	$[\text{Rn}]5f^{14}6d^{10}7s^27p^2$	-42691.493680
Br	$[\text{Ar}]3d^{10}4s^24p^5$	-2572.270918	Re	$[\text{Xe}]4f^{14}5d^7$	-15783.943765	Mc	$[\text{Rn}]5f^{14}6d^{10}7s^27p^3$	-43583.961779
Kr	$[\text{Ar}]3d^{10}4s^24p^6$	-2752.054977	Os	$[\text{Xe}]4f^{14}5d^8$	-16290.259414	Lv	$[\text{Rn}]5f^{14}6d^{10}7s^27p^4$	-44486.902550
Rb	$[\text{Kr}]5s^1$	-2938.319660	Ir	$[\text{Xe}]4f^{14}5d^9$	-16805.965623	Ts	$[\text{Rn}]5f^{14}6d^{10}7s^27p^5$	-45400.354767
Sr	$[\text{Kr}]5s^2$	-3131.545686	Pt	$[\text{Xe}]4f^{14}5d^{10}$	-17331.121868	Og	$[\text{Rn}]5f^{14}6d^{10}7s^27p^6$	-46324.355815
Y	$[\text{Kr}]5s^25p^1$	-3331.559557	Au	$[\text{Xe}]4f^{14}5d^{10}6s^1$	-17865.342083			
Zr	$[\text{Kr}]5s^24d^2$	-3538.662298	Hg	$[\text{Xe}]4f^{14}5d^{10}6s^2$	-18408.991495			

and implementation purposes. For instance, Clementi-Roetti wave functions [135] are often used for non-self-consistent benchmarks of density functionals, but the availability of a program for self-consistent calculations is certain to help future developments as numerical instabilities in a functional may not be detected in non-self-consistent calculations.

Furthermore, we have reported the nonrelativistic spin-restricted ground-state configurations of all atoms in the periodic table at HF and HFS levels of theory. Such knowledge is useful for implementations of the superposition of atomic densities guess [136,137], which is often implemented based on spin-restricted fractionally occupied calculations.

TABLE V. Nonrelativistic spin-restricted spherical HFS configurations for all elements in the periodic table. Entries in italics indicate that a lower-lying configuration was identified but it failed to converge.

H	$1s^1$	-0.406534	Nb	$[\text{Kr}]4d^35s^2$	-3747.428127	Tl	$[\text{Xe}]4f^{14}5d^{10}6s^26p^1$	-18948.496862
He	$1s^2$	-7.223640	Mo	$[\text{Kr}]5s^14d^5$	-3969.125868	Pb	$[\text{Xe}]4f^{14}5d^{10}6s^26p^2$	-19510.422489
Li	$[\text{He}]2s^1$	-7.174881	Tc	$[\text{Kr}]4d^65s^1$	-4198.246878	Bi	$[\text{Xe}]4f^{14}5d^{10}6s^26p^3$	-20081.732046
Be	$[\text{He}]2s^2$	-14.223291	Ru	$[\text{Kr}]4d^8$	-4434.888516	Po	$[\text{Xe}]4f^{14}5d^{10}6s^26p^4$	-20662.460965
B	$[\text{He}]2s^22p^1$	-24.050406	Rh	$[\text{Kr}]4d^9$	-4679.115070	At	$[\text{Xe}]4f^{14}5d^{10}6s^26p^5$	-21252.645251
C	$[\text{He}]2s^22p^2$	-37.053605	Pd	$[\text{Kr}]4d^{10}$	-4931.010033	Rn	$[\text{Xe}]4f^{14}5d^{10}6s^26p^6$	-21852.321426
N	$[\text{He}]2s^22p^3$	-53.567903	Ag	$[\text{Kr}]4d^{10}5s^1$	-5190.567420	Fr	$[\text{Rn}]7s^1$	-22461.201212
O	$[\text{He}]2s^22p^4$	-73.925425	Cd	$[\text{Kr}]4d^{10}5s^2$	-5457.821825	Ra	$[\text{Rn}]7s^2$	-23079.470637
F	$[\text{He}]2s^22p^5$	-98.456607	In	$[\text{Kr}]4d^{10}5s^25p^1$	-5732.640932	Ac	$[\text{Rn}]7s^25f^1$	-23707.189388
Ne	$[\text{He}]2s^22p^6$	-127.490741	Sn	$[\text{Kr}]4d^{10}5s^25p^2$	-6015.182678	Th	$[\text{Rn}]5f^27s^2$	-24344.622650
Na	$[\text{Ne}]3s^1$	-160.628228	Sb	$[\text{Kr}]4d^{10}5s^25p^3$	-6305.500906	Pa	$[\text{Rn}]5f^37s^2$	-24991.833379
Mg	$[\text{Ne}]3s^2$	-198.248792	Te	$[\text{Kr}]4d^{10}5s^25p^4$	-6603.649656	U	$[\text{Rn}]7s^15f^5$	-25648.893676
Al	$[\text{Ne}]3s^23p^1$	-240.346857	I	$[\text{Kr}]4d^{10}5s^25p^5$	-6909.683446	Np	$[\text{Rn}]7s^15f^6$	-26315.863733
Si	$[\text{Ne}]3s^23p^2$	-287.145287	Xe	$[\text{Kr}]4d^{10}5s^25p^6$	-7223.657213	Pu	$[\text{Rn}]7s^15f^7$	-26992.780160
P	$[\text{Ne}]3s^23p^3$	-338.804261	Cs	$[\text{Xe}]6s^1$	-7545.272707	Am	$[\text{Rn}]5f^87s^1$	-27679.697021
S	$[\text{Ne}]3s^23p^4$	-395.481609	Ba	$[\text{Xe}]6s^2$	-7874.734118	Cm	$[\text{Rn}]5f^97s^1$	-28376.667807
Cl	$[\text{Ne}]3s^23p^5$	-457.333996	La	$[\text{Xe}]6s^24f^1$	-8212.148603	Bk	$[\text{Rn}]5f^{10}7s^1$	-29083.745568
Ar	$[\text{Ne}]3s^23p^6$	-524.517426	Ce	$[\text{Xe}]6s^24f^2$	-8557.852692	Cf	$[\text{Rn}]5f^{11}7s^1$	-29800.983007
K	$[\text{Ar}]4s^1$	-596.699051	Pr	$[\text{Xe}]4f^36s^2$	-8911.927706	Es	$[\text{Rn}]5f^{12}7s^1$	-30528.432552
Ca	$[\text{Ar}]4s^2$	-674.160118	Nd	$[\text{Xe}]4f^46s^2$	-9274.451612	Fm	$[\text{Rn}]5f^{13}7s^1$	-31266.146407
Sc	$[\text{Ar}]4s^23d^1$	-757.000629	Pm	$[\text{Xe}]4f^56s^2$	-9645.500832	Md	$[\text{Rn}]5f^{14}7s^1$	-32014.176598
Ti	$[\text{Ar}]4s^23d^2$	-845.497930	Sm	$[\text{Xe}]4f^66s^2$	-10025.150892	No	$[\text{Rn}]5f^{14}7s^2$	-32772.269829
V	$[\text{Ar}]3d^34s^2$	-939.796100	Eu	$[\text{Xe}]4f^76s^2$	-10413.476735	Lr	$[\text{Rn}]5f^{14}7s^26d^1$	-33540.454380
Cr	$[\text{Ar}]3d^44s^2$	-1040.034946	Gd	$[\text{Xe}]4f^86s^2$	-10810.552897	Rf	$[\text{Rn}]5f^{14}6d^27s^2$	-34318.854809
Mn	$[\text{Ar}]4s^13d^6$	-1146.366756	Tb	$[\text{Xe}]4f^96s^2$	-11216.453617	Db	$[\text{Rn}]5f^{14}6d^47s^1$	-35107.525943
Fe	$[\text{Ar}]4s^13d^7$	-1258.917212	Dy	$[\text{Xe}]4f^{10}6s^2$	-11631.252911	Sg	$[\text{Rn}]5f^{14}6d^6$	-35906.506548
Co	$[\text{Ar}]4s^13d^8$	-1377.819755	Ho	$[\text{Xe}]4f^{11}6s^2$	-12055.024619	Bh	$[\text{Rn}]5f^{14}6d^7$	-36715.824635
Ni	$[\text{Ar}]3d^94s^1$	-1503.210775	Er	$[\text{Xe}]4f^{12}6s^2$	-12487.842443	Hs	$[\text{Rn}]5f^{14}6d^8$	-37535.505151
Cu	$[\text{Ar}]3d^{10}4s^1$	-1635.226377	Tm	$[\text{Xe}]4f^{13}6s^2$	-12929.779972	Mt	$[\text{Rn}]5f^{14}6d^9$	-38365.584348
Zn	$[\text{Ar}]3d^{10}4s^2$	-1773.909886	Yb	$[\text{Xe}]4f^{14}6s^2$	-13380.910702	Ds	$[\text{Rn}]5f^{14}6d^{10}$	-39206.098757
Ga	$[\text{Ar}]3d^{10}4s^24p^1$	-1919.085911	Lu	$[\text{Xe}]4f^{14}6s^25d^1$	-13840.976253	Rg	$[\text{Rn}]5f^{14}6d^{10}7s^1$	-40056.951158
Ge	$[\text{Ar}]3d^{10}4s^24p^2$	-2070.946515	Hf	$[\text{Xe}]4f^{14}6s^25d^2$	-14310.121254	Cn	$[\text{Rn}]5f^{14}6d^{10}7s^2$	-40918.195130
As	$[\text{Ar}]3d^{10}4s^24p^3$	-2229.571620	Ta	$[\text{Xe}]4f^{14}6s^25d^3$	-14788.392156	Nh	$[\text{Rn}]5f^{14}6d^{10}7s^27p^1$	-41789.700671
Se	$[\text{Ar}]3d^{10}4s^24p^4$	-2395.043625	W	$[\text{Xe}]4f^{14}6s^15d^5$	-15275.846800	Fl	$[\text{Rn}]5f^{14}6d^{10}7s^27p^2$	-42671.589032
Br	$[\text{Ar}]3d^{10}4s^24p^5$	-2567.446685	Re	$[\text{Xe}]4f^{14}5d^66s^1$	-15772.541265	Mc	$[\text{Rn}]5f^{14}6d^{10}7s^27p^3$	-43563.886976
Kr	$[\text{Ar}]3d^{10}4s^24p^6$	-2746.866101	Os	$[\text{Xe}]4f^{14}5d^8$	-16278.531177	Lv	$[\text{Rn}]5f^{14}6d^{10}7s^27p^4$	-44466.621119
Rb	$[\text{Kr}]5s^1$	-2932.972209	Ir	$[\text{Xe}]4f^{14}5d^9$	-16793.845129	Ts	$[\text{Rn}]5f^{14}6d^{10}7s^27p^5$	-45379.818244
Sr	$[\text{Kr}]5s^2$	-3125.998090	Pt	$[\text{Xe}]4f^{14}5d^{10}$	-17318.533845	Og	$[\text{Rn}]5f^{14}6d^{10}7s^27p^6$	-46303.505356
Y	$[\text{Kr}]5s^24d^1$	-3325.964742	Au	$[\text{Xe}]4f^{14}5d^{10}6s^1$	-17852.550237			
Zr	$[\text{Kr}]5s^24d^2$	-3533.076869	Hg	$[\text{Xe}]4f^{14}5d^{10}6s^2$	-18395.920112			

The present approach is useful for implementations of the SAP guess [79]. For instance, the implementation of SAP now available in the development version of the PSI4 program [138] is based on HFS potentials tabulated during the present work. Instead of the 4000 point tabulation used in [79] with unknown error, the ten-element calculations of the present work yield 751-point tabulations that repro-

duce the sub-micro-Hartree-level accuracy of the original calculation.

The atomic orbitals obtained from the present approach may also be useful for initializing fully numerical molecular electronic structure calculations via either a superposition of atomic densities or in combination to the extended Hückel rule developed in [79].

ACKNOWLEDGMENTS

I thank Dirk Andrae, Volker Blum, Eli Kraissler, Jacek Kobus, Leeor Kronik, Micael Oliveira, Dage Sundholm, Edward Valeev, and Lucas Visscher for discussions.

Computational resources provided by CSC—It Center for Science Ltd. (Espoo, Finland) and the Finnish Grid and Cloud Infrastructure are gratefully acknowledged. This work has been supported by the Academy of Finland (Suomen Akatemia) through Project No. 311149.

- [1] S. Lehtola, A review on non-relativistic, fully numerical electronic structure calculations on atoms and diatomic molecules, *Int. J. Quantum Chem.* **119**, e25968 (2019).
- [2] D. R. Hartree, W. Hartree, and B. Swirles, Self-consistent field, including exchange and superposition of configurations, with some results for oxygen, *Philos. Trans. R. Soc. London A* **238**, 229 (1939).
- [3] P. Hohenberg and W. Kohn, Inhomogeneous electron gas, *Phys. Rev.* **136**, B864 (1964).
- [4] W. Kohn and L. J. Sham, Self-consistent equations including exchange and correlation effects, *Phys. Rev.* **140**, A1133 (1965).
- [5] E. J. Baerends, V. Branchadell, and M. Sodupe, Atomic reference energies for density functional calculations, *Chem. Phys. Lett.* **265**, 481 (1997).
- [6] P. J. Hay, Gaussian basis sets for molecular calculations. The representation of 3d orbitals in transition-metal atoms, *J. Chem. Phys.* **66**, 4377 (1977).
- [7] R. F. Prat, Stability of self-consistent symmetries in atomic theory: The cases of Ne, F^- , and O^- , *Phys. Rev. A* **6**, 1735 (1972).
- [8] A. Goursot, J. P. Malrieu, and D. R. Salahub, Bonding in C_2 and Be_2 : Broken symmetry and correlation in DFT solutions, *Theor. Chim. Acta* **91**, 225 (1995).
- [9] F. W. Kutzler and G. S. Painter, Energies of Atoms with Non-spherical Charge Densities Calculated with Nonlocal Density-Functional Theory, *Phys. Rev. Lett.* **59**, 1285 (1987).
- [10] P. H. T. Philipsen and E. J. Baerends, Cohesive energy of 3d transition metals: Density functional theory atomic and bulk calculations, *Phys. Rev. B* **54**, 5326 (1996).
- [11] J. Tao and J. P. Perdew, Nonempirical Construction of Current-Density Functionals from Conventional Density-Functional Approximations, *Phys. Rev. Lett.* **95**, 196403 (2005).
- [12] E. R. Johnson, R. M. Dickson, and A. D. Becke, Density functionals and transition-metal atoms, *J. Chem. Phys.* **126**, 184104 (2007).
- [13] S. Pittalis, S. Kurth, and E. K. U. Gross, On the degeneracy of atomic states within exact-exchange (spin-) density functional theory, *J. Chem. Phys.* **125**, 084105 (2006).
- [14] S. Lehtola, Fully numerical Hartree–Fock and density functional calculations. I. Atoms, *Int. J. Quantum Chem.* **119**, e25945 (2019).
- [15] L. N. Anderson, M. Belén Oviedo, and Bryan M. Wong, Accurate electron affinities and orbital energies of anions from a nonempirically tuned range-separated density functional theory approach, *J. Chem. Theory Comput.* **13**, 1656 (2017).
- [16] A. Unsöld, Beiträge zur quantenmechanik der atome, *Ann. Phys.* **387**, 355 (1927).
- [17] M. Levy, Electron densities in search of Hamiltonians, *Phys. Rev. A* **26**, 1200 (1982).
- [18] H. Englisch and R. Englisch, Exact density functionals for ground-state energies. I. General results, *Phys. Status Solidi* **123**, 711 (1984).
- [19] H. Englisch and R. Englisch, Exact density functionals for ground-state energies II. Details and remarks, *Phys. Status Solidi* **124**, 373 (1984).
- [20] B. I. Dunlap and W. N. Mei, Basis set effects on spectroscopic constants for C_2 and Si_2 and the symmetry dilemma in the $X\alpha$ model, *J. Chem. Phys.* **78**, 4997 (1983).
- [21] S. G. Wang and W. H. E. Schwarz, Simulation of non-dynamical correlation in density functional calculations by the optimized fractional orbital occupation approach: Application to the potential energy surfaces of O_3 and SO_2 , *J. Chem. Phys.* **105**, 4641 (1996).
- [22] P. R. T. Schipper, O. V. Gritsenko, and E. J. Baerends, One-determinantal pure state versus ensemble Kohn–Sham solutions in the case of strong electron correlation: CH_2 and C_2 , *Theor. Chem. Acc.* **99**, 329 (1998).
- [23] R. Takeda, S. Yamanaka, and K. Yamaguchi, Fractional occupation numbers and spin density functional calculations of degenerate systems, *Int. J. Quantum Chem.* **93**, 317 (2003).
- [24] C. R. Nygaard and J. Olsen, The energy, orbitals and electric properties of the ozone molecule with ensemble density functional theory, *Mol. Phys.* **111**, 1259 (2013).
- [25] M. M. Valiev and G. W. Fernando, Occupation numbers in density-functional calculations, *Phys. Rev. B* **52**, 10697 (1995).
- [26] K. J. H. Giesbertz and E. J. Baerends, Aufbau derived from a unified treatment of occupation numbers in Hartree–Fock, Kohn–Sham, and natural orbital theories with the Karush–Kuhn–Tucker conditions for the inequality constraints $n_i \leq 1$ and $n_i \geq 0$, *J. Chem. Phys.* **132**, 194108 (2010).
- [27] E. Kraissler, G. Makov, N. Argaman, and I. Kelson, Fractional occupation in Kohn–Sham density-functional theory and the treatment of non-pure-state ν -representable densities, *Phys. Rev. A* **80**, 032115 (2009).
- [28] N. D. Mermin, Thermal properties of the inhomogeneous electron gas, *Phys. Rev.* **137**, A1441 (1965).
- [29] M. V. Stoitsov and I. Zh. Petkov, Density functional theory at finite temperatures, *Ann. Phys. (NY)* **184**, 121 (1988).
- [30] J.-D. Chai, Density functional theory with fractional orbital occupations, *J. Chem. Phys.* **136**, 154104 (2012).
- [31] J.-D. Chai, Thermally-assisted-occupation density functional theory with generalized-gradient approximations, *J. Chem. Phys.* **140**, 18A521 (2014).
- [32] C.-S. Wu and J.-D. Chai, Electronic properties of zigzag graphene nanoribbons studied by TAO-DFT, *J. Chem. Theory Comput.* **11**, 2003 (2015).
- [33] C.-N. Yeh and J.-D. Chai, Role of Kekulé and non-Kekulé structures in the radical character of alternant polycyclic

- aromatic hydrocarbons: A TAO-DFT study, *Sci. Rep.* **6**, 30562 (2016).
- [34] S. Seenithurai and J.-D. Chai, Effect of Li adsorption on the electronic and hydrogen storage properties of acenes: A dispersion-corrected TAO-DFT study, *Sci. Rep.* **6**, 33081 (2016).
- [35] C.-S. Wu, P.-Y. Lee, and J.-D. Chai, Electronic properties of cyclacenes from TAO-DFT, *Sci. Rep.* **6**, 37249 (2016).
- [36] J.-D. Chai, Role of exact exchange in thermally-assisted-occupation density functional theory: A proposal of new hybrid schemes, *J. Chem. Phys.* **146**, 044102 (2017).
- [37] S. Seenithurai and J.-D. Chai, Effect of Li termination on the electronic and hydrogen storage properties of linear carbon chains: A TAO-DFT study, *Sci. Rep.* **7**, 4966 (2017).
- [38] C.-Y. Lin, K. Hui, J.-H. Chung, and J.-D. Chai, Self-consistent determination of the fictitious temperature in thermally-assisted-occupation density functional theory, *RSC Adv.* **7**, 50496 (2017).
- [39] C.-N. Yeh, C. Wu, H. Su, and J.-D. Chai, Electronic properties of the coronene series from thermally-assisted-occupation density functional theory, *RSC Adv.* **8**, 34350 (2018).
- [40] S. Seenithurai and J.-D. Chai, Electronic and hydrogen storage properties of Li-terminated linear boron chains studied by TAO-DFT, *Sci. Rep.* **8**, 13538 (2018).
- [41] Q. Deng and J.-D. Chai, Electronic properties of triangle-shaped graphene nanoflakes from TAO-DFT, *ACS Omega* **4**, 14202 (2019).
- [42] J.-H. Chung and J.-D. Chai, Electronic properties of Möbius cyclacenes studied by thermally-assisted-occupation density functional theory, *Sci. Rep.* **9**, 2907 (2019).
- [43] S. Grimme, Towards first principles calculation of electron impact mass spectra of molecules, *Angew. Chemie Int. Ed.* **52**, 6306 (2013).
- [44] S. Grimme and A. Hansen, A practicable real-space measure and visualization of static electron-correlation effects, *Angew. Chemie Int. Ed.* **54**, 12308 (2015).
- [45] A. D. Rabuck and G. E. Scuseria, Improving self-consistent field convergence by varying occupation numbers, *J. Chem. Phys.* **110**, 695 (1999).
- [46] P. Kratzer and J. Neugebauer, The basics of electronic structure theory for periodic systems, *Front. Chem.* **7**, 1 (2019).
- [47] P. Slavíček and T. J. Martínez, Ab initio floating occupation molecular orbital-complete active space configuration interaction: An efficient approximation to CASSCF, *J. Chem. Phys.* **132**, 234102 (2010).
- [48] C. L. Fu and K. M. Ho, First-principles calculation of the equilibrium ground-state properties of transition metals: Applications to Nb and Mo, *Phys. Rev. B* **28**, 5480 (1983).
- [49] M. Methfessel and A. T. Paxton, High-precision sampling for Brillouin-zone integration in metals, *Phys. Rev. B* **40**, 3616 (1989).
- [50] N. Marzari, D. Vanderbilt, A. De Vita, and M. C. Payne, Thermal Contraction and Disordering of the Al(110) Surface, *Phys. Rev. Lett.* **82**, 3296 (1999).
- [51] J. M. Holender, M. J. Gillan, M. C. Payne, and A. D. Simpson, Static, dynamic, and electronic properties of liquid gallium studied by first-principles simulation, *Phys. Rev. B* **52**, 967 (1995).
- [52] M. Springborg, R. C. Albers, and K. Schmidt, Fractional occupancies and temperature in electronic-structure calculations, *Phys. Rev. B* **57**, 1427 (1998).
- [53] O. Grotheer and M. Fähnle, Correction terms to the density-functional ground-state energy arising from occupation number broadening, *Phys. Rev. B* **58**, 13459 (1998).
- [54] M. J. Mehl, Occupation-number broadening schemes: Choice of “temperature”, *Phys. Rev. B* **61**, 1654 (2000).
- [55] M. Weinert and J. W. Davenport, Fractional occupations and density-functional energies and forces, *Phys. Rev. B* **45**, 13709 (1992).
- [56] R. W. Warren and B. I. Dunlap, Fractional occupation numbers and density functional energy gradients within the linear combination of Gaussian-type orbitals approach, *Chem. Phys. Lett.* **262**, 384 (1996).
- [57] P. Pulay, Convergence acceleration of iterative sequences. The case of SCF iteration, *Chem. Phys. Lett.* **73**, 393 (1980).
- [58] P. Pulay, Improved SCF convergence acceleration, *J. Comput. Chem.* **3**, 556 (1982).
- [59] F. W. Averill and G. S. Painter, Steepest-descent determination of occupation numbers and energy minimization in the local-density approximation, *Phys. Rev. B* **46**, 2498 (1992).
- [60] E. Cancès, K. N. Kudin, G. E. Scuseria, and G. Turinici, Quadratically convergent algorithm for fractional occupation numbers in density functional theory, *J. Chem. Phys.* **118**, 5364 (2003).
- [61] C. R. Nygaard and J. Olsen, A second-order unconstrained optimization method for canonical-ensemble density-functional methods, *J. Chem. Phys.* **138**, 094109 (2013).
- [62] M. J. Gillan, Calculation of the vacancy formation energy in aluminium, *J. Phys.: Condens. Matter* **1**, 689 (1989).
- [63] G. W. Fernando, G.-X. Qian, M. Weinert, and J. W. Davenport, First-principles molecular dynamics for metals, *Phys. Rev. B* **40**, 7985 (1989).
- [64] M. P. Grumbach, D. Hohl, R. M. Martin, and R. Car, *Ab initio* molecular dynamics with a finite-temperature density functional, *J. Phys.: Condens. Matter* **6**, 1999 (1994).
- [65] N. Chetty, M. Weinert, T. S. Rahman, and J. W. Davenport, Vacancies and impurities in aluminum and magnesium, *Phys. Rev. B* **52**, 6313 (1995).
- [66] N. Marzari, D. Vanderbilt, and M. C. Payne, Ensemble Density-Functional Theory for *Ab Initio* Molecular Dynamics of Metals and Finite-Temperature Insulators, *Phys. Rev. Lett.* **79**, 1337 (1997).
- [67] S. Kotochigova, Z. H. Levine, E. L. Shirley, M. D. Stiles, and C. W. Clark, Local-density-functional calculations of the energy of atoms, *Phys. Rev. A* **55**, 191 (1997).
- [68] I.-H. Lee and R. M. Martin, Applications of the generalized-gradient approximation to atoms, clusters, and solids, *Phys. Rev. B* **56**, 7197 (1997).
- [69] J. P. Perdew, K. Burke, and M. Ernzerhof, Generalized Gradient Approximation Made Simple, *Phys. Rev. Lett.* **77**, 3865 (1996).
- [70] J. P. Perdew, K. Burke, and M. Ernzerhof, Generalized Gradient Approximation Made Simple [Phys. Rev. Lett. 77, 3865 (1996)], *Phys. Rev. Lett.* **78**, 1396 (1997).
- [71] M. Fuchs and M. Scheffler, *Ab initio* pseudopotentials for electronic structure calculations of poly-atomic systems using

- density-functional theory, *Comput. Phys. Commun.* **119**, 67 (1999).
- [72] M. J. T. Oliveira and F. Nogueira, Generating relativistic pseudo-potentials with explicit incorporation of semi-core states using APE, the Atomic Pseudo-potentials Engine, *Comput. Phys. Commun.* **178**, 524 (2008).
- [73] T. Ozaki and H. Kino, Numerical atomic basis orbitals from H to Kr, *Phys. Rev. B* **69**, 195113 (2004).
- [74] V. Blum, R. Gehrke, F. Hanke, P. Havu, V. Havu, X. Ren, K. Reuter, and M. Scheffler, *Ab initio* molecular simulations with numeric atom-centered orbitals, *Comput. Phys. Commun.* **180**, 2175 (2009).
- [75] J. Andzelm, E. Radzio, and D. R. Salahub, Compact basis sets for LCAO-LSD calculations. Part I: Method and bases for Sc to Zn, *J. Comput. Chem.* **6**, 520 (1985).
- [76] N. Godbout, D. R. Salahub, J. Andzelm, and E. Wimmer, Optimization of Gaussian-type basis sets for local spin density functional calculations. Part I. Boron through neon, optimization technique and validation, *Can. J. Chem.* **70**, 560 (1992).
- [77] D. Porezag and M. Pederson, Optimization of Gaussian basis sets for density-functional calculations, *Phys. Rev. A* **60**, 2840 (1999).
- [78] G. te Velde, F. M. Bickelhaupt, E. J. Baerends, C. Fonseca Guerra, S. J. A. van Gisbergen, J. G. Snijders, and T. Ziegler, Chemistry with ADF, *J. Comput. Chem.* **22**, 931 (2001).
- [79] S. Lehtola, Assessment of initial guesses for self-consistent field calculations. Superposition of atomic potentials: Simple yet efficient, *J. Chem. Theory Comput.* **15**, 1593 (2019).
- [80] J. C. Slater, J. B. Mann, T. M. Wilson, and J. H. Wood, Non-integral occupation numbers in transition atoms in crystals, *Phys. Rev.* **184**, 672 (1969).
- [81] S. F. Abdunur, J. Linderberg, U. Öhrn, and P. W. Thulstrup, Atomic central-field models for open shells with application to transition metals, *Phys. Rev. A* **6**, 889 (1972).
- [82] J. Janak, Proof that $\partial E/\partial n_i = \epsilon_i$ in density-functional theory, *Phys. Rev. B* **18**, 7165 (1978).
- [83] E. Kraisler, G. Makov, and I. Kelson, Ensemble v -representable *ab initio* density-functional calculation of energy and spin in atoms: A test of exchange-correlation approximations, *Phys. Rev. A* **82**, 042516 (2010).
- [84] S. Lehtola, HelFEM—Finite element methods for electronic structure calculations on small systems, see <https://github.com/susilehtola/HelFEM>.
- [85] S. Lehtola, Fully numerical Hartree-Fock and density functional calculations. II. Diatomic molecules, *Int. J. Quantum Chem.* **119**, e25944 (2019).
- [86] P. E. Blöchl, Projector augmented-wave method, *Phys. Rev. B* **50**, 17953 (1994).
- [87] P. Schwerdtfeger, The pseudopotential approximation in electronic structure theory, *Chem. Phys. Chem.* **12**, 3143 (2011).
- [88] J. R. Trail and R. J. Needs, Norm-conserving Hartree-Fock pseudopotentials and their asymptotic behavior, *J. Chem. Phys.* **122**, 014112 (2005).
- [89] W. A. Al-Saidi, E. J. Walter, and A. M. Rappe, Optimized norm-conserving Hartree-Fock pseudopotentials for plane-wave calculations, *Phys. Rev. B* **77**, 075112 (2008).
- [90] X. Wu, E. J. Walter, A. M. Rappe, R. Car, and A. Selloni, Hybrid density functional calculations of the band gap of $\text{Ga}_x\text{In}_{1-x}\text{N}$, *Phys. Rev. B* **80**, 115201 (2009).
- [91] J. Yang, L. Z. Tan, and A. M. Rappe, Hybrid functional pseudopotentials, *Phys. Rev. B* **97**, 085130 (2018).
- [92] J. C. Slater, The theory of complex spectra, *Phys. Rev.* **34**, 1293 (1929).
- [93] S. Lehtola, C. Steigemann, M. J. T. Oliveira, and M. A. L. Marques, Recent developments in LIBXC—A comprehensive library of functionals for density functional theory, *SoftwareX* **7**, 1 (2018).
- [94] P. Verma and R. J. Bartlett, Increasing the applicability of density functional theory. IV. Consequences of ionization-potential improved exchange-correlation potentials, *J. Chem. Phys.* **140**, 18A534 (2014).
- [95] Y. Jin and R. J. Bartlett, The QTP family of consistent functionals and potentials in Kohn-Sham density functional theory, *J. Chem. Phys.* **145**, 034107 (2016).
- [96] R. L. A. Haiduke and R. J. Bartlett, Non-empirical exchange-correlation parametrizations based on exact conditions from correlated orbital theory, *J. Chem. Phys.* **148**, 184106 (2018).
- [97] R. Peverati and D. G. Truhlar, Screened-exchange density functionals with broad accuracy for chemistry and solid-state physics, *Phys. Chem. Chem. Phys.* **14**, 16187 (2012).
- [98] P. Verma, Y. Wang, S. Ghosh, X. He, and D. G. Truhlar, Revised M11 exchange-correlation functional for electronic excitation energies and ground-state properties, *J. Phys. Chem. A* **123**, 2966 (2019).
- [99] N. Mardirossian and M. Head-Gordon, ω B97X—V: A 10-parameter, range-separated hybrid, generalized gradient approximation density functional with nonlocal correlation, designed by a survival-of-the-fittest strategy, *Phys. Chem. Chem. Phys.* **16**, 9904 (2014).
- [100] N. Mardirossian and M. Head-Gordon, ω B97M—V: A combinatorially optimized, range-separated hybrid, meta-GGA density functional with VV10 nonlocal correlation, *J. Chem. Phys.* **144**, 214110 (2016).
- [101] J. G. Ángyán, I. Gerber, and M. Marsman, Spherical harmonic expansion of short-range screened Coulomb interactions, *J. Phys. A* **39**, 8613 (2006).
- [102] F. Zapata, E. Luppi, and J. Toulouse, Linear-response range-separated density-functional theory for atomic photoexcitation and photoionization spectra, *J. Chem. Phys.* **150**, 234104 (2019).
- [103] P. M. W. Gill, R. D. Adamson, and J. A. Pople, Coulomb-attenuated exchange energy density functionals, *Mol. Phys.* **88**, 1005 (1996).
- [104] T. Leininger, H. Stoll, H.-J. Werner, and A. Savin, Combining long-range configuration interaction with short-range density functionals, *Chem. Phys. Lett.* **275**, 151 (1997).
- [105] T. Yanai, D. P. Tew, and N. C. Handy, A new hybrid exchange-correlation functional using the Coulomb-attenuating method (CAM-B3LYP), *Chem. Phys. Lett.* **393**, 51 (2004).
- [106] H. Yukawa, On the interaction of elementary particles. I, *Proc. Phys. Math. Soc. Jpn., 3rd Ser.* **17**, 48 (1935).
- [107] R. J. Harrison, G. I. Fann, T. Yanai, Z. Gan, and G. Beylkin, Multiresolution quantum chemistry: Basic theory and initial applications, *J. Chem. Phys.* **121**, 11587 (2004).
- [108] L. Frediani, E. Fossgaard, T. Flå, and K. Ruud, Fully adaptive algorithms for multivariate integral equations using the non-standard form and multiwavelets with applications to the Poisson and bound-state Helmholtz kernels in three dimensions, *Mol. Phys.* **111**, 1143 (2013).

- [109] F. Jensen, Unifying general and segmented contracted basis sets. Segmented polarization consistent basis sets, *J. Chem. Theory Comput.* **10**, 1074 (2014).
- [110] E. Solala, S. A. Losilla, D. Sundholm, W. Xu, and P. Parkkinen, Optimization of numerical orbitals using the Helmholtz kernel, *J. Chem. Phys.* **146**, 084102 (2017).
- [111] P. Parkkinen, S. A. Losilla, E. Solala, E. A. Toivanen, W.-H. Xu, and D. Sundholm, A generalized grid-based fast multipole method for integrating Helmholtz Kernels, *J. Chem. Theory Comput.* **13**, 654 (2017).
- [112] M. Seth and T. Ziegler, Range-separated exchange functionals with Slater-type functions, *J. Chem. Theory Comput.* **8**, 901 (2012).
- [113] M. Seth, T. Ziegler, M. Steinmetz, and S. Grimme, Modeling transition metal reactions with range-separated functionals, *J. Chem. Theory Comput.* **9**, 2286 (2013).
- [114] J. F. Rico, R. López, G. Ramírez, and I. Ema, Repulsion integrals involving Slater-type functions and Yukawa potential, *Theor. Chem. Acc.* **132**, 1304 (2013).
- [115] Y. Akinaga and S. Ten-no, Range-separation by the Yukawa potential in long-range corrected density functional theory with Gaussian-type basis functions, *Chem. Phys. Lett.* **462**, 348 (2008).
- [116] S. Ten-no, Initiation of explicitly correlated Slater-type geminal theory, *Chem. Phys. Lett.* **398**, 56 (2004).
- [117] S. Ten-no, New implementation of second-order Møller-Plesset perturbation theory with an analytic Slater-type geminal, *J. Chem. Phys.* **126**, 014108 (2007).
- [118] Y. Akinaga and S. Ten-No, Intramolecular charge-transfer excitation energies from range-separated hybrid functionals using the Yukawa potential, *Int. J. Quantum Chem.* **109**, 1905 (2009).
- [119] J. Heyd, G. E. Scuseria, and M. Ernzerhof, Hybrid functionals based on a screened Coulomb potential, *J. Chem. Phys.* **118**, 8207 (2003).
- [120] R. Ahlrichs, A simple algebraic derivation of the Obara-Saika scheme for general two-electron interaction potentials, *Phys. Chem. Chem. Phys.* **8**, 3072 (2006).
- [121] S. L. Marshall, Calculation of Coulombic lattice potentials: II. Spherical harmonic expansion of the Green function, *J. Phys.: Condens. Matter* **14**, 3175 (2002).
- [122] J. A. Pople, P. M. W. Gill, and B. G. Johnson, Kohn-Sham density-functional theory within a finite basis set, *Chem. Phys. Lett.* **199**, 557 (1992).
- [123] F. Della Sala, E. Fabiano, and L. A. Constantin, Kinetic-energy-density dependent semilocal exchange-correlation functionals, *Int. J. Quantum Chem.* **116**, 1641 (2016).
- [124] T. Kato, On the eigenfunctions of many-particle systems in quantum mechanics, *Commun. Pure Appl. Math.* **10**, 151 (1957).
- [125] E. Steiner, Charge densities in atoms, *J. Chem. Phys.* **39**, 2365 (1963).
- [126] Á. Nagy and K. D. Sen, Ground- and excited-state cusp conditions for the electron density, *J. Chem. Phys.* **115**, 6300 (2001).
- [127] R. Sharp and G. Horton, A variational approach to the unipotent many-electron problem, *Phys. Rev.* **90**, 317 (1953).
- [128] J. P. Perdew and Y. Wang, Accurate and simple analytic representation of the electron-gas correlation energy, *Phys. Rev. B* **45**, 13244 (1992).
- [129] Eli Kraisler (private communication).
- [130] H. Iikura, T. Tsuneda, T. Yanai, and K. Hirao, A long-range correction scheme for generalized-gradient-approximation exchange functionals, *J. Chem. Phys.* **115**, 3540 (2001).
- [131] C. Lee, W. Yang, and R. G. Parr, Development of the Colle-Salvetti correlation-energy formula into a functional of the electron density, *Phys. Rev. B* **37**, 785 (1988).
- [132] J. Lehtola, M. Hakala, A. Sakko, and K. Hämäläinen, ERKALE—A flexible program package for X-ray properties of atoms and molecules, *J. Comput. Chem.* **33**, 1572 (2012).
- [133] A. D. Becke, A new mixing of Hartree-Fock and local density-functional theories, *J. Chem. Phys.* **98**, 1372 (1993).
- [134] S. L. Saito, Hartree-Fock-Roothaan energies and expectation values for the neutral atoms He to Uuo: The B-spline expansion method, *At. Data Nucl. Data Tables* **95**, 836 (2009).
- [135] E. Clementi and C. Roetti, Roothaan-Hartree-Fock atomic wave functions, *At. Data Nucl. Data Tables* **14**, 177 (1974).
- [136] J. Almlöf, K. Faegri, and K. Korsell, Principles for a direct SCF approach to LCAO-MO ab initio calculations, *J. Comput. Chem.* **3**, 385 (1982).
- [137] J. H. Van Lenthe, R. Zwaans, H. J. J. Van Dam, and M. F. Guest, Starting SCF calculations by superposition of atomic densities, *J. Comput. Chem.* **27**, 926 (2006).
- [138] R. M. Parrish, L. A. Burns, D. G. A. Smith, A. C. Simmonett, A. E. DePrince, E. G. Hohenstein, U. Bozkaya, A. Yu. Sokolov, R. Di Remigio, R. M. Richard, J. F. Gonthier, A. M. James, H. R. McAlexander, A. Kumar, M. Saitow, X. Wang, B. P. Pritchard, P. Verma, H. F. Schaefer, K. Patkowski, R. A. King, E. F. Valeev, F. A. Evangelista, J. M. Turney, T. D. Crawford, and C. D. Sherrill, Psi4 1.1: An open-source electronic structure program emphasizing automation, advanced libraries, and interoperability, *J. Chem. Theory Comput.* **13**, 3185 (2017).