



Making ab initio QED functional(s): Nonperturbative and photon-free effective frameworks for strong light–matter coupling

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Strong light–matter coupling provides a promising path for the control of quantum matter where the latter is routinely described from first principles. However, combining the quantized nature of light with this ab initio tool set is challenging and merely developing as the coupled light–matter Hilbert space is conceptually different and computational cost quickly becomes overwhelming. In this work, we provide a nonperturbative photon-free formulation of quantum electrodynamics (QED) in the long-wavelength limit, which is formulated solely on the matter Hilbert space and can serve as an accurate starting point for such ab initio methods. The present formulation is an extension of quantum mechanics that recovers the exact results of QED for the zero- and infinite-coupling limit and the infinite-frequency as well as the homogeneous limit, and we can constructively increase its accuracy. We show how this formulation can be used to devise approximations for quantum-electrodynamical density-functional theory (QEDFT), which in turn also allows us to extend the ansatz to the full minimal-coupling problem and to nonadiabatic situations. Finally, we provide a simple local density–type functional that takes the strong coupling to the transverse photon degrees of freedom into account and includes the correct frequency and polarization dependence. This QEDFT functional accounts for the quantized nature of light while remaining computationally simple enough to allow its application to a large range of systems. All approximations allow the seamless application to periodic systems.

quantum-electrodynamical density-functional theory (QEDFT) | cavity quantum electrodynamics | strong light–matter coupling | photon-free QED | QEDFT functionals

In the last decade, seminal experimental results (1–4) have demonstrated that the properties and dynamics of atoms, molecules, and solids can be substantially modified by coupling strongly to the modes of a photonic environment. The strong coupling between light and matter in these cases leads to the emergence of hybrid light–matter states (polaritons) which subsequently can be used to control, for instance, chemical reactions (5–11) and enhance charge and energy transport (12–19), even over very large distances, and there are recent indications that they might be used to increase the critical temperature of superconductors (20–22). Many of these changes persist at normal ambient conditions and are hence promising for quantum-technological applications (1). They question, however, our common perception that light and matter are distinct physical entities and call for a more unified description.

Typically, in such strong light–matter coupling situations, only a few photonic modes, which are supported by a cavity geometry, play a substantial role. Their interaction with matter is, however, greatly enhanced in comparison to the free-space situation. Effective subwavelength confinement (2, 23–26) or circuit

geometries (27–29) enable hybridization strengths on the order of the matter excitation. A noticeable hybridization between light and matter (strong coupling) results in the emergence of polaritons in the excited states. The coupling between (artificial) cavity modes and matter excitation can be so considerable that all eigenvalues of the individual constituents are affected, and even the ground state becomes correlated (ultrastrong coupling). For even larger hybridization, the character of light and matter becomes truly interlacing (deep ultrastrong coupling).

Many of the experimentally observed effects, in particular those involving chemical reactions under strong coupling, are so far theoretically not well understood (6, 11, 30–32). Detailed theoretical explanations are largely missing as strong light–matter coupling calls in principle for investigations within the framework of quantum electrodynamics (QED) (33). This, however, leads to an enormous increase of computational complexity as the combined light–matter Hilbert space becomes prohibitively large (illustrated in Fig. 1).

The most common approach is to considerably reduce the dimensionality of the light–matter Hilbert space by deciding a

Significance

Strong interaction between light and matter recently attracted enormous interest after seminal work highlighted how materials and chemical reactions can be effectively controlled with cavities, suggesting new quantum technological applications and alternative paths to catalysis. Combining the quantized nature of light and matter is, however, a nontrivial task due to the overwhelming complexity of the combined system. We first motivate a nonperturbative photon-free reformulation of quantum electrodynamics (QED) that recovers, even in its simplest form, the correct physical limits of the original QED problem, providing us with an efficient framework for further developments. By combination with density-functional theory, we obtain the first nonperturbative local density–type functional that describes the quantized interaction between light and matter while remaining computationally feasible.

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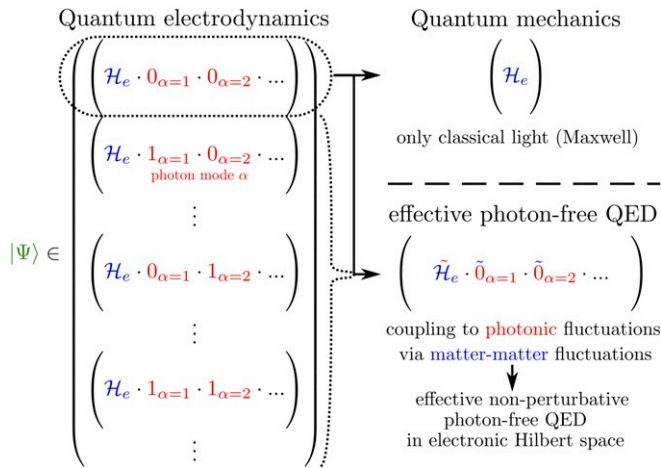


Fig. 1. An illustration of the exponential increase in computational complexity: In QED the matter Hilbert space \mathcal{H}_e is extended with the photonic Fock space (indicated by the vectors for different modes α starting with the vacuum state $0_{\alpha=1} \cdot 0_{\alpha=2} \cdot \dots$), and hence, the combined Hilbert space (Left) grows exponentially also with the number of photonic states. A vast reduction of complexity is found by approximating QED with quantum mechanics, where the photonic sector is subsumed into the effective mass of the particles and into classical electromagnetic fields. The effective photon-free QED ansatz (Bottom Right) takes the dominant fluctuations of the quantized electromagnetic field $\hat{0}$ into account, while having the same dimensionality as ordinary quantum mechanics (Top Right).

priori which matter and photon states are supposed to be important. This leads to effective model light–matter Hamiltonians (3, 34). A different approach is to reformulate the full QED problem in terms of reduced quantities. Quantum mechanics, for instance, can be viewed as a reduction of the full QED problem onto the matter sector only, where the photon field is taken into account approximately by the effective (physical) mass of the particles (33, 35) and by the longitudinal Coulomb interaction as well as by possible Van der Waals corrections (33, 35, 36). This can be viewed as an example of an effective ab initio light–matter Hamiltonian. This simplification is, however, no longer valid in the case of strong light–matter coupling, and more advanced reformulations of QED become necessary. An in principle exact reformulation of QED on the basis of reduced quantities is quantum-electrodynamical density-functional theory (QEDFT), which allows one to avoid the unfeasible coupled matter–photon wave function altogether (37–40). The main drawback of QEDFT is that we need to find approximations to the matter–photon coupling, usually in terms of matter quantities only (41, 42). Deriving such functional expressions for interactions is already for matter-only DFT a challenging task, and the photonic field introduces additional and unfamiliar components into the problem. It therefore becomes desirable to define an effective ab initio photon-free QED as an alternative to standard quantum mechanics which is also applicable to strong light–matter coupling situations and serves as a starting point for approximations for QEDFT. Furthermore, this effective matter-only theory should be constructively improvable and recover the physical limits of the original QED solution.

In this work we provide such a nonperturbative photon-free QED reformulation that takes the dominant fluctuations of the quantized electromagnetic field explicitly into account (Section 1). It thereby lifts the artificial distinction between light and matter that underlies standard quantum mechanics, and it remains applicable from the weak to the deep ultrastrong coupling regime. Among other things we show that the ansatz recovers the

exact QED solution in the weak- and infinite-coupling limit and the infinite-frequency limit, as well as for the homogeneous electron gas, without the need to treat the photonic degrees of freedom at all. While many effective ab initio descriptions of light–matter coupling problems are known, e.g., the high-frequency limit of Floquet theory (43–45), this high consistency with fundamental physical conditions provides a much more flexible and general perspective. Since the theory is built upon an explicit ansatz for photonic operators in terms of matter quantities, we still have approximate access to photonic observables. Further, we show that the nonperturbative photon-free QED reformulation is obtained as a truncation of an especially efficient basis expansion of the full QED problem (Section 2). In this way, we can consistently increase the accuracy of the ansatz, converging to the original QED results and accounting for all correlations between light and matter. Furthermore, we show how this ansatz can be used to set up an orbital-dependent approximation in QEDFT that shares the same beneficial features as the nonperturbative photon-free QED reformulation (Section 3). Based on this functional we propose a simple local density–type approximation for strong light–matter coupling that provides up to ultrastrong coupling excellent results and yet accounts for the correct frequency and cavity polarization dependence. We benchmark all introduced approximations against the exact solution of one-dimensional hydrogen coupled to a single cavity mode and further benchmarks for a double-well potential can be found in *SI Appendix*. Our derivations can be applied seamlessly to periodic systems and many mode cavities. Due to the flexibility of the QEDFT framework, we can finally discuss how this ansatz can be extended to full minimal coupling and comment on nonadiabatic extensions.

1. Photon-Free QED Ansatz

The point of departure for our endeavor is the Pauli–Fierz (or minimal coupling) Hamiltonian of nonrelativistic QED in Coulomb gauge. For simplicity, we will focus here at first on the long-wavelength limit, but the following conceptions are general enough to allow an abstraction beyond this common simplification. We provide an outlook beyond those common approximations in Section 3. In atomic units with electron charge $q = -|e| = -1$ and keeping the nuclei fixed we have

$$\hat{H}_{\text{PF}} = \frac{1}{2} \sum_{i=1}^{N_e} \left(-i\nabla_i + \frac{1}{c} \hat{\mathbf{A}} \right)^2 + \sum_{i=1}^{N_e} v(\mathbf{r}_i) + \frac{1}{2} \sum_{i \neq j}^{N_e} w(\mathbf{r}_i, \mathbf{r}_j) + \sum_{\alpha=1}^{M_p} \omega_{\alpha} \left(\hat{a}_{\alpha}^{\dagger} \hat{a}_{\alpha} + \frac{1}{2} \right). \quad [1]$$

The N_e electrons move according to the forces caused by the local nuclear potentials v in addition to their longitudinal interaction w and the coupling to the transverse vector potential at the molecular center-of-charge \mathbf{r}_0

$$\hat{\mathbf{A}} = \sqrt{4\pi c^2} \sum_{\alpha=1}^{M_p} S_{\alpha}(\mathbf{r}_0) \epsilon_{\alpha} \frac{1}{\sqrt{2\omega_{\alpha}}} \left(\hat{a}_{\alpha}^{\dagger} + \hat{a}_{\alpha} \right). \quad [2]$$

The M_p cavity eigenmodes $S_{\alpha}(\mathbf{r}_0) \propto 1/\sqrt{V}$ are the solutions to the boundary value problem of the effective cavity (which we assume to be lossless for simplicity) and provide the local (vacuum) field strength. The vector potential is conveniently expanded in the eigenmodes of the cavity geometry $\hat{\mathbf{A}} = \sum_{\alpha} \hat{A}_{\alpha} \epsilon_{\alpha}$, featuring polarization ϵ_{α} , frequency ω_{α} , and cavity volume V . The \hat{A}_{α} obey the following equation of motion in the

Heisenberg picture,

$$\left(\frac{1}{c^2} \frac{d^2}{dt^2} + \frac{\omega_\alpha^2}{c^2}\right) \hat{A}_{\alpha,H} = -\frac{\lambda_\alpha^2}{c} \epsilon_\alpha \cdot (\hat{\mathbf{J}}_{p,H} - \hat{\mathbf{J}}_{d,H}), \quad [3]$$

with $\lambda_\alpha = \sqrt{4\pi} S_\alpha(\mathbf{r}_0)$ the fundamental coupling strength. The paramagnetic current $\hat{\mathbf{J}}_p = -i \sum_i \nabla_i$ serves as a driving force for the photonic degrees of freedom. The diamagnetic current in the long-wavelength approximation $\hat{\mathbf{J}}_d = -N_e/c \sum_\alpha \hat{A}_\alpha \epsilon_\alpha$, on the other hand, can be conveniently absorbed in effective cavity frequencies and polarizations denoted from here on as $\tilde{\omega}_\alpha$ and $\tilde{\epsilon}_\alpha$. For a single mode, we can simply move $\hat{\mathbf{J}}_d$ to the left-hand side to obtain $\tilde{\omega}_\alpha^2 = \omega_\alpha^2 + \omega_{d,\alpha}^2$, $\omega_{d,\alpha}^2 = N_e \lambda_\alpha^2$, and $\epsilon_\alpha = \tilde{\epsilon}_\alpha$. The diamagnetic term therefore dresses the bare cavity frequency by the fundamental coupling strength λ_α and the amount of charged particles. For multiple modes, a normal-mode (or Bogoliubov) transformation (Section 5.1) similarly eliminates the diamagnetic current, leading to a pure bilinear coupling in the Hamiltonian Eq. 1. The representation in normal modes with $\tilde{\omega}_\alpha$ also transforms all related operators \hat{A} where we keep the original notation for brevity.

After absorbing the diamagnetic current, the Ehrenfest, or more specifically Maxwell's, equation associated to Eq. 3 is easily solved with the help of the classical Green's function for the harmonic oscillator,

$$A_\alpha(t) = -\tilde{\omega}_\alpha \int_{-\infty}^t c \frac{\omega_{d,\alpha}^2}{N_e \tilde{\omega}_\alpha^2} \sin(\tilde{\omega}_\alpha(t-t')) \tilde{\epsilon}_\alpha \cdot \mathbf{J}_p(t') dt'. \quad [4]$$

In the limit of Maxwell's equation there is therefore no need to keep track of the photonic degrees of freedom, as their evolution is fully determined by the initial conditions and the matter degrees of freedom.

Now let us stretch this idea, comparably to the relativistic Breit substitution (46), to the Heisenberg equation of motion, and let us attempt to avoid the necessity of photonic degrees of freedom in Eq. 1. Certainly, such a substitution is far less trivial for the operator quantity; it is furthermore complicated by the fact that now we deal with two different pictures, i.e., Heisenberg for Eq. 3 and Schrödinger for Eq. 1. A suitable first approximation is an adiabatic ansatz for the quantum fluctuations,

$$\Delta \hat{A}_\alpha \approx -c \frac{\omega_{d,\alpha}^2}{N_e \tilde{\omega}_\alpha^2} \tilde{\epsilon}_\alpha \cdot \Delta \hat{\mathbf{J}}_p \text{ where } \Delta \hat{O} = \hat{O} - \langle \hat{O} \rangle. \quad [5]$$

Surely, also the pure photonic contributions have to be substituted, and we obtain

$$\tilde{\omega}_\alpha \hat{a}_\alpha^\dagger \hat{a}_\alpha \approx \frac{1}{2} \frac{\omega_{d,\alpha}^2}{N_e \tilde{\omega}_\alpha^2} (\tilde{\epsilon}_\alpha \cdot \Delta \hat{\mathbf{J}}_p) (\tilde{\epsilon}_\alpha \cdot \hat{\mathbf{J}}_p) \quad [6]$$

in the adiabatic limit (Section 5.2). Replacing now all photonic degrees of freedom by their introduced counterparts, we obtain the effective photon-free Hamiltonian

$$\begin{aligned} \hat{H}_{\text{pof}}(t) = & -\frac{1}{2} \sum_{i=1}^{N_e} \nabla_i^2 + \sum_{i=1}^{N_e} v(\mathbf{r}_i) + \frac{1}{2} \sum_{i \neq j}^{N_e} w(\mathbf{r}_i, \mathbf{r}_j) \\ & + \sum_{\alpha=1}^{M_p} \frac{\tilde{\omega}_\alpha}{2} - \sum_{\alpha=1}^{M_p} \frac{\omega_{d,\alpha}^2}{N_e \tilde{\omega}_\alpha^2} \left[\frac{1}{2} \tilde{\epsilon}_\alpha \cdot (\hat{\mathbf{J}}_p - \mathbf{J}_p(t)) \right. \\ & \left. + \tilde{\omega}_\alpha \int_{-\infty}^t \sin(\tilde{\omega}_\alpha(t-t')) \tilde{\epsilon}_\alpha \cdot \mathbf{J}_p(t') dt' \right] (\tilde{\epsilon}_\alpha \cdot \hat{\mathbf{J}}_p), \end{aligned} \quad [7]$$

which is now time-dependent. While computationally simpler to solve, more importantly, this provides us with a starting point

conceptually much closer to the known realm of electronic structure theory. Although the photon-free Hamiltonian depends on time, all equilibrium solutions are time-translational invariant, meaning that they depend on the excitation energy which labels a given excited state but not on time. Excited states can be calculated using linear response theory.

A particularly interesting feature of this effective Hamiltonian Eq. 7 is that the adiabatic correction takes the form of a polarization-projected kinetic operator $-(\tilde{\epsilon}_\alpha \cdot \hat{\mathbf{J}}_p)^2 \sim (\tilde{\epsilon}_\alpha \cdot \nabla_i)^2 + (\tilde{\epsilon}_\alpha \cdot \nabla_i)(\tilde{\epsilon}_\alpha \cdot \nabla_{j \neq i})$ with opposite sign to the kinetic operator. A large part of the photonic fluctuations are therefore responsible for dynamically increasing the mass of the charged particle along the axis of polarization, very much in line with the perturbative mass renormalization obtained from the Lamb shift (35) and the accumulation of electronic density inside cavities (19, 36, 42). If we extend the number of modes to infinity, we even recover the logarithmic ultraviolet divergence characteristic of the Lamb shift (47). We notice furthermore that simply disregarding pure photonic contributions in the substitution procedure from Eqs. 1–7 would mean to miss the factor $\frac{1}{2}$ introduced in Eq. 6. This would imply that for $\lambda_\alpha \rightarrow \infty$ the mass would tend not to infinity but to negative values instead, and the Hamiltonian would become unbounded from below, a common problem for similar approaches within relativistic regimes (48).

While we accomplished our initial goal of a nonperturbative photon-free QED theory, such an adiabatic substitution does not come without sacrifices. Fig. 2 illustrates the absorption spectrum of one-dimensional hydrogen strongly coupled to an optical cavity using the exact Pauli–Fierz Hamiltonian, the photon-free description in Eq. 7, and the purely classical description of the photonic field. Clearly, even on this strongly simplified level including only adiabatic fluctuations, the photon-free description in Eq. 7 improves noticeably over the semiclassical Maxwell–Schrödinger picture. It accounts correctly for the upward (first matter excitation) and downward (higher matter excitations) bending at low cavity frequencies. The description of multiphoton excitations present in the exact solution would demand to go beyond the adiabatic approximation in Eq. 5.

A particularly important feature of the photon-free Hamiltonian is the explicit dependence on the cavity frequency via $\omega_{d,\alpha}^2/\tilde{\omega}_\alpha^2$. To understand the importance of this observation, let us utilize the Power–Zienau–Woolley (PZW) transformation (49–51) and rewrite our original Coulomb-gauge Hamiltonian Eq. 1 in the routinely utilized PZW form (33, 35, 51)

$$\begin{aligned} \hat{H}_{\text{PZW}} = & -\frac{1}{2} \sum_{i=1}^{N_e} \nabla_i^2 + \sum_{i=1}^{N_e} v(\mathbf{r}_i) + \frac{1}{2} \sum_{i \neq j}^{N_e} w(\mathbf{r}_i, \mathbf{r}_j) \\ & + \frac{1}{2} \sum_{\alpha=1}^{M_p} \left[(-i\partial_{p_\alpha})^2 + \omega_\alpha^2 \left(p_\alpha + \frac{\lambda_\alpha}{\omega_\alpha} \tilde{\epsilon}_\alpha \cdot \sum_{i=1}^{N_e} \mathbf{r}_i \right)^2 \right]. \end{aligned} \quad [8]$$

The photonic harmonic-oscillator coordinates are given here explicitly as p_α . Without truncation of electronic or photonic space, both the Pauli–Fierz Eq. 1 and the PZW–Hamiltonian Eq. 8 provide the exact same result for all physical observables, as we would expect from the concept of gauge invariance. We note, however, that the PZW Hamiltonian is not a convenient starting point for extended periodic systems, since the periodicity in the matter coordinates is explicitly broken (52). Eq. 8 now involves the dipole self-polarization term $\frac{1}{2} (\lambda_\alpha \tilde{\epsilon}_\alpha \cdot \sum_i \mathbf{r}_i)^2$. So even if we disregard the photonic degrees of freedom altogether, we remain with a confining harmonic potential acting on the electronic degrees of freedom that results in the correct behavior for zero and infinite coupling. This approximation is sometimes referred to as QED Hartree–Fock (36, 53). Fig. 3,

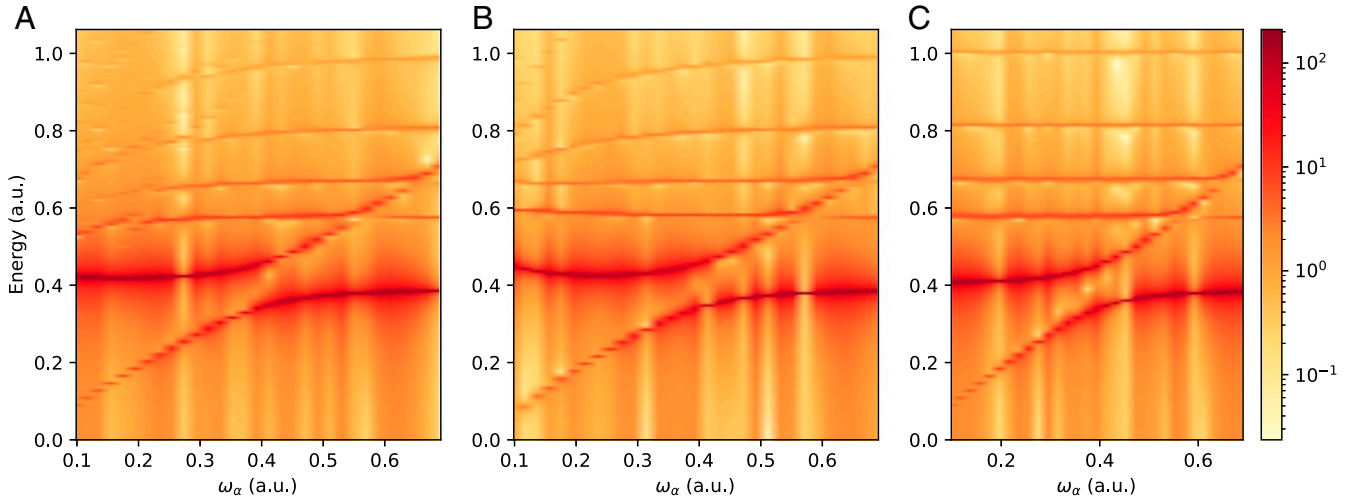


Fig. 2. Linear-response spectrum of the dipole moment $|r(\omega_\alpha)|$ for a single cavity mode with different cavity frequencies ω_α coupled to one-dimensional soft-Coulomb hydrogen $v(x) = -1/\sqrt{x^2+1}$. Shown are (A) the exact reference, (B) the photon-free effective Hamiltonian Eq. 7, and (C) the semiclassical Maxwell-Schrödinger solution. The fundamental coupling strength was chosen such that $g_\alpha/\omega_\alpha = 0.136$, $g_\alpha = \sqrt{\omega_\alpha/2}\lambda_{\alpha r}$, i.e., on the interface between strong and ultrastrong coupling, with $\omega_\alpha = \varepsilon_1^{\text{hydrogen}} - \varepsilon_0^{\text{hydrogen}}$ in resonance with the first electronic excitation. We have chosen a grid of 301 points with $0.1 a_0$ spacing (a_0 being the Bohr radius) and 40 photonic Fock states for the exact reference using the PZW Hamiltonian Eq. 8. The response was obtained by the delta-kick method applying a perturbation $v_{\text{kick}}(x, t) = -10^{-4}/\pi \cdot 10^{-2}/[(t-1)^2 + 10^{-4}]x$ with consecutive time propagation for $T = 10^3$ atomic units (a.u.) with fourth-order Runge-Kutta and a time-stepping of $dt = 5 \cdot 10^{-4}$ a.u. Vertical lines are artifacts originating from numerical deviations, logarithmic scale, and interpolation.

however, clearly illustrates that the photon-free Hamiltonian Eq. 7 is superior due to its explicit frequency dependence. The latter furthermore guarantees that photon-free and exact description coincide for infinite cavity frequency, where matter excitations are no longer allowed to couple to the photonic degrees of freedom. It provides thus an excellent electronic subspace solution which becomes exact whenever a factorized wave function ansatz is possible.

By handling operators as if we would treat expectation values, we intrinsically assume that both sides of Eq. 5 possess the same set of eigenstates. This, for instance, is satisfied for a set of plane waves describing free electrons and quantized photonic fields. Unfortunately, for the vast majority of condensed matter systems this condition will not be met. Knowing about the limitation of such a construction, we can, however, use the solution of the homogeneous electron gas inside the cavity as a basis for a formally more accurate description. We show in the following that even by restricting ourselves to the original electronic Hilbert space this provides accurate results which become exact in the limit of weak and deep ultrastrong coupling, thus providing a preferable starting point for functional development.

2. The Photon-Coupled Homogeneous Electron Gas Basis

We already stated that the photon-free ansatz becomes exact in the homogeneous limit. This is indeed no coincidence, as it can be shown that in the free ($v=0$) and noninteracting ($w=0$) limit, the Pauli-Fierz Hamiltonian Eq. 1 can be diagonalized analytically with a combination of Bogoliubov and coherent shift transformations (47). The obtained photon-coupled homogeneous electron gas (pHEG) Hamiltonian is purely additive in the new transformed operators,

$$\hat{H}_{\text{pHEG}} = -\frac{1}{2} \sum_{i=1}^{N_e} \nabla_i^2 - \sum_{\alpha=1}^{M_p} \tilde{\omega}_\alpha \hat{\beta}_\alpha^2 + \sum_{\alpha=1}^{M_p} \tilde{\omega}_\alpha \left(\hat{c}_\alpha^\dagger \hat{c}_\alpha + \frac{1}{2} \right), \quad [9]$$

with the coherent shift operator

$$\hat{\beta}_\alpha = \frac{\omega_{d,\alpha}}{\sqrt{2N_e}\tilde{\omega}_\alpha^3} \tilde{\epsilon}_\alpha \cdot \sum_{i=1}^{N_e} (-i\nabla_i).$$

The precise definition of the transformed annihilation and creation operators $\hat{c}_\alpha, \hat{c}_\alpha^\dagger$ can be found in *SI Appendix*. An immediate consequence of the additive structure is that the eigenfunctions of the Hamiltonian Eq. 9 are factorized in nature. Its eigenbasis can be chosen as plane-wave Slater determinants and displaced photon number states,

$$|\{\mathbf{k}_j\}, \{\beta_\alpha(\mathbf{K}), n_\alpha\}\rangle = |\Phi_{\{\mathbf{k}_j\}}\rangle \otimes \prod_{\alpha=1}^{M_p} |\beta_\alpha(\mathbf{K}), n_\alpha\rangle.$$

Here $\mathbf{K} = \sum_i \mathbf{k}_i$ represents the collective momentum of all particles, a well-defined quantum number in the pHEG system. The photonic states $|\beta_\alpha(\mathbf{K}), n_\alpha\rangle$ implicitly account for the collective momenta of the homogeneous electronic system. It becomes apparent that this new basis, although factorized, represents intrinsically interacting light and matter parts, very similar to the photon-free ansatz motivated in Section 1. It is now our intention to bring this particular pHEG solution into use as a basis for a general, inhomogeneous system. This demands an expression of the local potential in the pHEG basis, a computationally simple task for any potential that can be represented in a Fourier basis. The matrix elements of the external single-particle potentials in the pHEG basis then read

$$\begin{aligned} & \langle \{\mathbf{k}_j\}, \{\beta_\alpha(\mathbf{K}), n_\alpha\} | v(\hat{\mathbf{r}}_i) | \{\mathbf{k}'_j\}, \{\beta_{\alpha'}(\mathbf{K}'), n'_{\alpha'}\} \rangle \quad [10] \\ & = \langle \Phi_{\{\mathbf{k}_j\}} | v(\hat{\mathbf{r}}_i) | \Phi_{\{\mathbf{k}'_j\}} \rangle \cdot \prod_{\alpha=1}^{M_p} \langle \beta_\alpha(\mathbf{K}), n_\alpha | \beta_{\alpha'}(\mathbf{K}'), n'_{\alpha'} \rangle, \end{aligned}$$

where all $\mathbf{k}_j = \mathbf{k}'_j$ have to match for $j \neq i$ in order to provide a nonzero expression due to orthogonality. The first part is the Fourier transformation of the potential at $\mathbf{k}_i - \mathbf{k}'_j$, and the second can be evaluated analytically in terms of associated Laguerre polynomials (54) with argument $\beta_\alpha(\mathbf{k}_i) - \beta_\alpha(\mathbf{k}'_j)$. We denote the

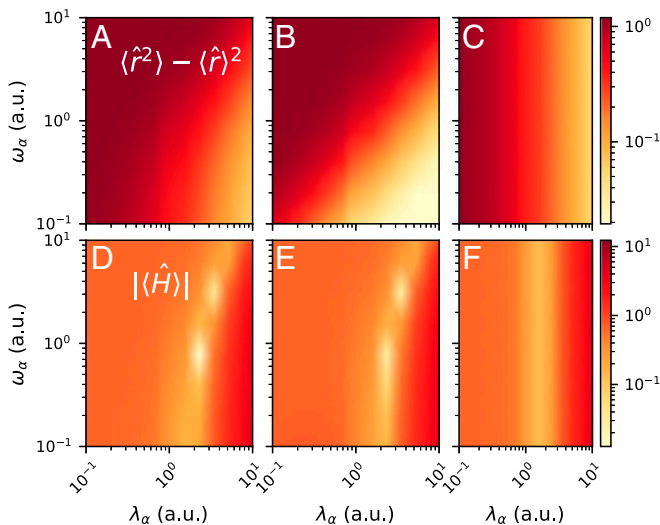


Fig. 3. Ground-state (A–C) variance and (D–F) energy for the one-dimensional soft-Coulomb hydrogen atom coupled to a single cavity mode. Illustrated are the exact solution (A and D), the photon-free Hamiltonian Eq. 7 (B and E), and the photon-free self-polarization solution of Eq. 8 (C and F). The results presented here are entirely due to the quantized nature of light; a semiclassical Maxwell–Schrödinger treatment would show no effect in the ground state. We disregarded the trivial zero-point energy shift $\omega_\alpha/2$ in all calculations and used 2,001 grid points with $0.025 a_0$ spacing, finite-difference order 4, and 40 photonic Fock states (using the PZW Hamiltonian).

second part as the Fourier transform of a function $m_\alpha^{n,n'}(\mathbf{r}_i)$, and consequently, Eq. 10 is conveniently expressed with the help of the convolution theorem,

$$\mathcal{F}v \cdot \prod_{\alpha=1}^{M_p} \mathcal{F}m_\alpha^{n,n'} = \mathcal{F} \left(v * \bigotimes_{\alpha=1}^{M_p} m_\alpha^{n,n'} \right).$$

The coherent shifts of the pHEG basis therefore provide an effective screening or mollification of the local potential. In the homogeneous system ($v=0$), photonic excitations n_α are mere replicas without substantial relevance. The moment our system becomes inhomogeneous ($v \neq 0$), however, we start to couple different eigenstates via scatterings at the local potential. In perfect agreement with our conclusions in Section 1, we observe that for an inhomogeneous system the photonic Fock space recovers its relevance. However, even in lowest order and therefore entirely within the zero-excitations sector, the modifier is nonzero,

$$m_\alpha^{0,0}(\mathbf{r}) = \sqrt{\frac{N_e \tilde{\omega}_\alpha^3}{\pi \omega_{d,\alpha}^2}} e^{-\frac{N_e \tilde{\omega}_\alpha^3}{\omega_{d,\alpha}^2} (\tilde{\epsilon}_\alpha \cdot \mathbf{r})^2} \delta(\tilde{\epsilon}_\alpha^{\perp,1} \cdot \mathbf{r}) \delta(\tilde{\epsilon}_\alpha^{\perp,2} \cdot \mathbf{r}),$$

where $\tilde{\epsilon}_\alpha^{\perp,1}, \tilde{\epsilon}_\alpha^{\perp,2}$ are selected such that they complete $\tilde{\epsilon}_\alpha$ into an orthonormal basis for \mathbb{R}^3 . Convolution with the Gaussian above acts as a mollification and yields a zeroth-order correction to our original photon-free ansatz Eq. 7. This correction vanishes, i.e., $m_\alpha^{0,0} \rightarrow \delta$, for $\lambda_\alpha \rightarrow 0$ or ∞ and $\omega_\alpha \rightarrow \infty$, and we observe again that the photon-free ansatz is exact in those limits. For higher orders n, n' the respectively modified potentials $v * m_\alpha^{n,n'}$ couple different excitation number sectors.

Assuming a sufficient number of photonic excitations n_α are considered, the pHEG basis provides exactly the same results as the original Pauli–Fierz Hamiltonian Eq. 1. As presented in Fig. 4, this Fock space dimension n_α can however be substantially smaller in the pHEG basis while still obtaining much better

converged energies than with the trivial noninteracting basis $|\Phi_{\{k_j\}}\rangle \otimes |n_\alpha\rangle$ for the Pauli–Fierz Hamiltonian. Especially in the ultrastrong coupling limit the superiority of the pHEG basis is apparent. Here the noninteracting basis demands a quickly growing Fock space while the pHEG basis is exact for $\lambda_\alpha \rightarrow \infty$ by construction, even in its lowest approximation. Very intuitively, the flatter the local potential, i.e., the closer our system resembles a homogeneous system, the better the pHEG basis converges. The sharper the local potential, the more scattering events have to be described by the pHEG basis. Comparing Fig. 4 A–D, it, however, becomes apparent that even for strongly localizing potentials this does not break the ansatz introduced here. Note that with the change of operators, photon observables are still accessible. Even in the photon-free case, matter fluctuations can represent photonic operators (see, e.g., Section 5.2). While we illustrate here the correlated energy, also purely photonic observables such as the photon number can be accurately reproduced by the truncated pHEG basis with very few excitations, as demonstrated in SI Appendix. The pHEG basis is similar to solving Hamiltonian Eq. 1 after a unitary transformation that moves the quantized fields into the spatial dependence of the external potential, similar to a Kramers–Henneberger transformation (55).

Despite the great advantage that the pHEG basis might provide, the high complexity of the electronic system remains. So even our nonperturbative photon-free ansatz will be challenged by any many-particle system and thus calls for further considerations if we intend to describe realistic molecules or solids. A combination of the photon-free ansatz with established electronic structure theory approaches will already provide a first suitable description of (ultra-) strong light–matter coupling. Section 3 illustrates how our previous considerations seamlessly integrate into QEDFT and shows how the conceptions behind the nonperturbative photon-free ansatz can be projected to realistic systems.

3. QEDFT

A much more reduced reformulation of QED is QEDFT, where the wave function of the coupled light–matter system is substituted by a current density and a vector potential (37–40, 56). Like in other versions of density-functional theory, no information is lost by this substitution, and the full wave function can be reconstructed in principle from this pair of reduced observables (37). The main drawback of QEDFT is, similar to any density-functional reformulation, that we in general do not have explicit expressions (expressed only by the pair of reduced observables) for the terms that appear in the defining equations (see, e.g., Eq. 13). To approximate these terms, one usually relies on auxiliary systems which are easier to treat numerically. The standard choice, which is also followed by and large in QEDFT, is to use noninteracting auxiliary systems. This Kohn–Sham construction gives rise to effective potentials that force the auxiliary noninteracting system to give the same current density and vector potential as the original reference system (37–40). These effective potentials can be expressed by differences of equations of motion. While it is relatively straightforward to find simple approximations for the longitudinal interaction (57, 58), for the transverse interactions between light and matter, there are only a few approximations hitherto available (41, 42, 59). The main problem with the matter–photon interaction terms in QEDFT is that the auxiliary, uncoupled Kohn–Sham wave function provides an inconvenient starting point to approximate a linearly coupled term of photonic and matter operators (see also the discussion of Eq. 13 below). It is at this point where QEDFT can strongly benefit from the methods devised here, and we can make use of the approximate theories derived above.

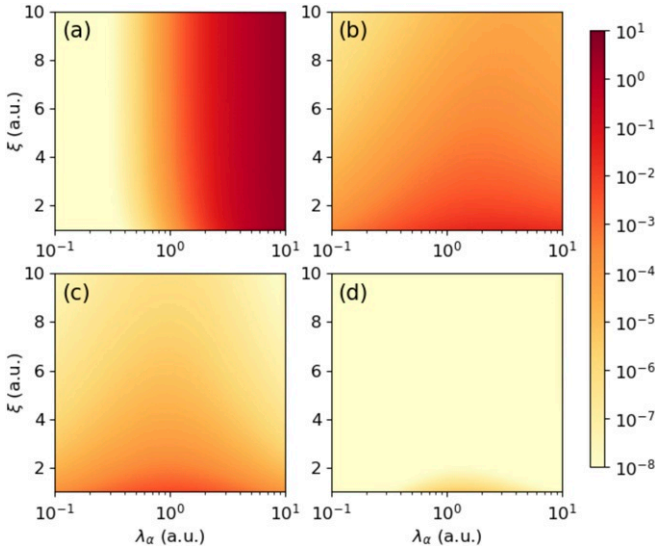


Fig. 4. Absolute deviation in ground-state energy compared to the exact Pauli-Fierz reference solution using $n_\alpha = 100$ photonic excitations of a tunable one-dimensional soft-Coulomb hydrogen $v(x) = -1/\sqrt{x^2 + \xi^2}$ coupled to a single cavity mode. (A) Pauli-Fierz Hamiltonian with max $n_\alpha = 4$ excitations, (B) pHEG basis with max $n_\alpha = 0$ and the original potential v , (C) pHEG basis with max $n_\alpha = 0$ but mollified potential $v * m_\alpha^{0,0}$, and (D) pHEG basis with max $n_\alpha = 4$. The pHEG basis is vastly superior over the noninteracting Pauli-Fierz basis for large λ_α . The flatter the local potential ($\xi \rightarrow \infty$), the closer we get to the scattering-free homogeneous solution and the better performs the pHEG basis. The electronic dimension was represented with 31 k points on a periodic grid.

Focusing on the ground-state case in the long-wavelength limit, the two basic equations of motion that we will use in order to define the effective potential are the balance of forces due to the paramagnetic current density $\hat{\mathbf{j}}_p(\mathbf{r}) = \frac{1}{2i} \sum_i (\delta(\mathbf{r} - \mathbf{r}_i) \vec{\nabla}_i - \vec{\nabla}_i \delta(\mathbf{r} - \mathbf{r}_i))$,

$$\rho(\mathbf{r}) \nabla v(\mathbf{r}) = \langle \hat{\mathbf{F}}_T(\mathbf{r}) \rangle + \langle \hat{\mathbf{F}}_W(\mathbf{r}) \rangle - \frac{1}{c} \langle (\hat{\mathbf{A}} \cdot \nabla) \hat{\mathbf{j}}_p(\mathbf{r}) \rangle, \quad [11]$$

together with the static mode-resolved Maxwell's equations,

$$A_\alpha = -\frac{c\omega_{d,\alpha}^2}{N_e \tilde{\omega}_\alpha^2} \tilde{\epsilon}_\alpha \cdot \mathbf{J}_p. \quad [12]$$

Both equations follow from the Heisenberg equation of motion for their respective operators (see also Eq. 3) with the Pauli-Fierz Hamiltonian of Eq. 1 and are evaluated with the ground state of the coupled light-matter system Ψ . Here the density operator is $\hat{\rho}(\mathbf{r}) = \sum_i \delta(\mathbf{r}_i - \mathbf{r})$, $\hat{\mathbf{F}}_T(\mathbf{r}) = \frac{i}{2} [\hat{\mathbf{j}}_p(\mathbf{r}), \sum_i \nabla_{r_i}^2]$ is the local stress force, $\hat{\mathbf{F}}_W(\mathbf{r}) = -\frac{i}{2} [\hat{\mathbf{j}}_p(\mathbf{r}), \sum_{i \neq j} w(\mathbf{r}_i, \mathbf{r}_j)]$ is the local interaction force, and $\langle \cdot \rangle$ indicates the expectation value with respect to Ψ . Alternatively, Eqs. 11 and 12 can be derived from the corresponding full minimal-coupling expression (SI Appendix) by taking the long-wavelength limit. In the static case the zero-component of the current density, i.e., the density $\rho(\mathbf{r})$, determines also the other components of the current density (39). Thus, the exact mean-field exchange-correlation (Mxc) scalar potential of static Kohn-Sham QEDFT is defined by

$$\begin{aligned} \nabla^2 v_{\text{Mxc}}(\mathbf{r}) = \nabla \cdot \frac{1}{\rho(\mathbf{r})} [\mathbf{F}_T([\Phi], \mathbf{r}) - \mathbf{F}_T([\Psi], \mathbf{r})] \\ - \mathbf{F}_W([\Psi], \mathbf{r}) + \frac{1}{c} \langle (\hat{\mathbf{A}} \cdot \nabla) \hat{\mathbf{j}}_p(\mathbf{r}) \rangle - \frac{1}{c} (\mathbf{A} \cdot \nabla) \mathbf{j}_p([\Phi], \mathbf{r}), \end{aligned} \quad [13]$$

where $\mathbf{F}_T([\Phi], \mathbf{r})$ and $\mathbf{j}_p([\Phi], \mathbf{r})$ indicate the expectation values of $\hat{\mathbf{F}}_T(\mathbf{r})$ and $\hat{\mathbf{j}}_p(\mathbf{r})$ with respect to the noninteracting Kohn-Sham wave function Φ , and $\mathbf{F}_T([\Psi], \mathbf{r})$ as well as $\mathbf{F}_W([\Psi], \mathbf{r})$ accordingly with respect to Ψ . Further, in the static case the mean-field contribution $(\mathbf{A} \cdot \nabla) \mathbf{j}_p[\Phi]$, which arises from the $\hat{\mathbf{J}}_p \cdot \mathbf{A}$ coupling in the Kohn-Sham system, is zero. For the differences in stress and interaction forces, various approximations based on the Kohn-Sham wave function exist in the DFT literature (60–63). Taking just the noninteracting Kohn-Sham wave function already leads to a nonvanishing expression for the interaction force $\mathbf{F}_W([\Phi], \mathbf{r})$. This contribution is called the local Hartree exchange contribution (58, 64),

$$\nabla^2 v_{\text{Hx}}(\mathbf{r}) = -\nabla \cdot \left[\frac{\mathbf{F}_W([\Phi], \mathbf{r})}{\rho(\mathbf{r})} \right]. \quad [14]$$

The same is no longer the case for the matter-photon interaction term in Eq. 13, where substituting just the auxiliary Kohn-Sham wave function leads to zero. The reason for this is that we work with an uncoupled and noninteracting Kohn-Sham wave function [the photonic part only consists of trivial shifted harmonic oscillators (39, 40)]. Having an approximation to the coupling in terms of matter quantities becomes therefore highly desirable.

The most straightforward approach follows the discussion in Section 1 by just replacing $\Delta \hat{A}_\alpha = \hat{A}_\alpha - A_\alpha \rightarrow -c\omega_{d,\alpha}^2 / (N_e \tilde{\omega}_\alpha^2) \tilde{\epsilon}_\alpha \cdot \Delta \hat{\mathbf{J}}_p$ in Eq. 13. In order to guarantee the real-valuedness of the expectation value we have to use a symmetrized form. While on the level of the equations of motion the validity of this ad hoc ansatz remains unclear, we find that by using the photon-free Hamiltonian of Eq. 7 to derive the equation of the paramagnetic current density the approximate coupling term becomes

$$\begin{aligned} \frac{1}{c} \langle (\Delta \hat{\mathbf{A}} \cdot \nabla) \hat{\mathbf{j}}_p(\mathbf{r}) \rangle \\ \rightarrow -\frac{1}{2} \sum_{\alpha=1}^{M_p} \frac{\omega_{d,\alpha}^2}{N_e \tilde{\omega}_\alpha^2} \left(\langle (\tilde{\epsilon}_\alpha \cdot \Delta \hat{\mathbf{J}}_p) (\tilde{\epsilon}_\alpha \cdot \nabla) \hat{\mathbf{j}}_p(\mathbf{r}) \rangle + c.c. \right), \end{aligned} \quad [15]$$

which exactly agrees with the above symmetrized equation of motion-based substitution. Here the expectation value is taken with respect to the photon-free QED ground-state wave function of Eq. 7. From this alternative approach that leads to the same expressions we see that this simple approximation becomes exact for the various limiting cases discussed in the previous sections. We furthermore know how to constructively increase the accuracy of this approach by including more terms from the pHEG basis expansion with the respective mollification. This implies that it is a reasonable strategy to build approximations for the coupling-force term based on the simple substitution of Eq. 15. Rewriting the exact Mxc potential as $v_{\text{Mxc}}(\mathbf{r}) = v_{\text{px}}(\mathbf{r}) + v_{\text{Hx}}(\mathbf{r}) + v_c(\mathbf{r})$, we find the static orbital-dependent photon exchange (px) contribution as

$$\nabla^2 v_{\text{px}}(\mathbf{r}) = -\nabla \cdot \left[\sum_{\alpha=1}^{M_p} \frac{\omega_{d,\alpha}^2}{N_e \tilde{\omega}_\alpha^2} \frac{(\tilde{\epsilon}_\alpha \cdot \nabla) \langle (\tilde{\epsilon}_\alpha \cdot \hat{\mathbf{J}}_p) \hat{\mathbf{j}}_p(\mathbf{r}) \rangle}{\rho(\mathbf{r})} \right], \quad [16]$$

where now the expectation value is taken with respect to a real-valued, auxiliary Kohn-Sham wave function and thus also $\mathbf{J}_p \equiv 0$. This expression can be further simplified for special cases, such

as in one spatial dimension and for one particle ($\Psi = \varphi = \sqrt{\rho}$), where it becomes

$$\begin{aligned} \partial_x^2 v_{\text{px}}(x) &= -\sum_{\alpha=1}^{M_p} \frac{\omega_{\text{d},\alpha}^2}{2\tilde{\omega}_{\alpha}^2} \partial_x \left[\frac{\partial_x [(\partial_x \varphi(x))^2 - (\partial_x^2 \varphi(x))\varphi(x)]}{\varphi(x)^2} \right] \\ &= \sum_{\alpha=1}^{M_p} \frac{\omega_{\text{d},\alpha}^2}{2\tilde{\omega}_{\alpha}^2} \partial_x^2 \left[\frac{\partial_x \sqrt{\rho(x)}}{\sqrt{\rho(x)}} \right]. \end{aligned} \quad [17]$$

Eq. 17 intuitively illustrates that electronic density becomes accumulated at local maxima, consistent with increasing the particle mass along the polarization direction as discussed in Section 1.

However, in general, the expression for v_{px} (Eq. 16) is still an orbital-dependent functional and hence can become costly for very large systems. It is therefore desirable to simplify this expression even further. Borrowing from a recent demonstration that Eq. 14 leads to the well-known exchange-only local-density approximation (LDA) for the Coulomb interaction in the homogeneous case (58), we follow the same strategy to devise a simple LDA for Eq. 16.

The starting point of the derivation, which we provide in detail in Section 5.3, is to express the current-current correlation in terms of one- and two-body reduced density matrices

$$\begin{aligned} &\langle (\tilde{\epsilon}_{\alpha} \cdot \hat{\mathbf{J}}_{\text{p}}) \hat{\mathbf{J}}_{\text{p}}(\mathbf{r}) \rangle \\ &= \frac{1}{2} [(\tilde{\epsilon}_{\alpha} \cdot \nabla') \nabla \rho_{(1)}(\mathbf{r}, \mathbf{r}') - (\tilde{\epsilon}_{\alpha} \cdot \nabla') \nabla' \rho_{(1)}(\mathbf{r}, \mathbf{r}')]_{\mathbf{r}'=\mathbf{r}} \\ &+ \int [(\tilde{\epsilon}_{\alpha} \cdot \nabla') \nabla \rho_{(2)}(\mathbf{r}, \mathbf{r}_2; \mathbf{r}', \mathbf{r}'_2) + c.c.]_{\mathbf{r}'=\mathbf{r}, \mathbf{r}'_2=\mathbf{r}_2} d\mathbf{r}_2. \end{aligned}$$

From here on, we use the closed-shell exchange representation of the two-body reduced density matrix $\rho_{(2)}(\mathbf{r}, \mathbf{r}_2; \mathbf{r}', \mathbf{r}'_2) = \frac{1}{2} [\rho_{(1)}(\mathbf{r}, \mathbf{r}') \rho_{(1)}(\mathbf{r}_2, \mathbf{r}'_2) - \frac{1}{2} \rho_{(1)}(\mathbf{r}, \mathbf{r}'_2) \rho_{(1)}(\mathbf{r}_2, \mathbf{r}')]_{\text{spin-}\frac{1}{2}}$ particles in combination with the homogeneous electron-gas ansatz $\rho_{(1)}(\mathbf{r}, \mathbf{r}') = 2(2\pi)^{-d} \int_{|\mathbf{k}| < k_{\text{F}}} \exp(i\mathbf{k} \cdot (\mathbf{r} - \mathbf{r}')) d\mathbf{k}$ with a local Fermi radius k_{F} in d spatial dimensions. This leads to the photon exchange-only LDA (pxLDA) that can be given for arbitrary spatial dimensionality d ,

$$\nabla^2 v_{\text{pxLDA}}(\mathbf{r}) = -\sum_{\alpha=1}^{M_p} \frac{2\pi^2 \omega_{\text{d},\alpha}^2}{N_e \tilde{\omega}_{\alpha}^2} (\tilde{\epsilon}_{\alpha} \cdot \nabla)^2 \left(\frac{\rho(\mathbf{r})}{2V_d} \right)^{\frac{2}{d}} \quad [18]$$

with V_d being the volume of the d -dimensional unit sphere. Let us emphasize that even this particularly simple functional includes the correct frequency and cavity-polarization dependence that are inherent to the nonperturbative photon-free approach. Assuming an isotropic interaction between light and matter, Eq. 18 allows for the direct solution $v_{\text{pxLDA}}(\mathbf{r}) = -\sum_{\alpha} 2\pi^2 \omega_{\text{d},\alpha}^2 / (dN_e \tilde{\omega}_{\alpha}^2) (\rho(\mathbf{r}) / (2V_d))^{\frac{2}{d}}$, a form suitable to describe free-space Lamb-shift physics. However, this form is consequently inapplicable to cavity settings.

The local potentials from Eqs. 13–18 are generally determined as the solutions to a Poisson-type equation $\nabla^2 v(\mathbf{r}) = -f(\mathbf{r})$. Solving the latter is a routinely performed calculation step in ab initio density-functional theory codes in order to obtain the Hartree potential $\nabla^2 v_{\text{H}}(\mathbf{r}) = -4\pi\rho(\mathbf{r})$ in a cost-efficient way (65). It is thus straightforward to go to realistic, three-dimensional systems. The numerical implementation and validation of such functionals for sizable, realistic systems will be discussed in detail in a forthcoming publication.

Finally, the energy associated with the $v_{\text{Mx}} = v_{\text{Hx}} + v_{\text{px}}$, or approximations thereof, is given by

$$\begin{aligned} E_{\text{Mx}} &= \left\langle -\frac{1}{2} \sum_{i=1}^{N_e} \nabla_i^2 + \frac{1}{2} \sum_{i \neq j}^{N_e} w(\mathbf{r}_i, \mathbf{r}_j) \right\rangle + \int v(\mathbf{r}) \rho(\mathbf{r}) d\mathbf{r} \\ &+ \sum_{\alpha=1}^{M_p} \frac{\omega_{\text{d},\alpha}^2}{2N_e \tilde{\omega}_{\alpha}^2} \left\langle \sum_{i,j=1}^{N_e} (\tilde{\epsilon}_{\alpha} \cdot \nabla_i) (\tilde{\epsilon}_{\alpha} \cdot \nabla_j) \right\rangle + \sum_{\alpha=1}^{M_p} \frac{\tilde{\omega}_{\alpha}}{2} \end{aligned}$$

with the expectation values with respect to the noninteracting Kohn–Sham wave function Φ determined by v_{Mx} . Further, photonic observables can be approximated by using the simple substitution discussed in Section 1 and also in Section 5.2.

Let us next consider how these simple exchange-type approximations for the coupling between light and matter perform in a test-case scenario. In Fig. 5 we show the spatial electronic dipole variance and total energy for a one-dimensional soft-Coulomb hydrogen model from weak to deep ultrastrong coupling. Investigating a single-particle system guarantees that we only consider the reliability of the light-matter approximation and do not mix in the approximate longitudinal electron-electron description. By construction, the photon exchange approximation v_{Mx} (blue diamonds) recovers the photon-free QED Hamiltonian solution (red dash-dotted line). Both (consistent) approximations overestimate the coupling effect (the exact result is the black dashed line) and tend to overbind. As discussed in Section 1, these approximations become exact for weak and deep ultrastrong coupling. In the intermediate (ultrastrong) regime, employing the mollification (crossed line and blue stars) suggested by the basis expansion in Section 2 leads to a clear improvement. The mollification also renders the minimization variational, i.e., by construction the energy is always above the exact energy. Similarly to the usual Coulombic LDA, which tends to delocalize electrons too strongly, the pxLDA also underestimates the enhanced binding. This leads to a partial error cancellation in the ultrastrong coupling domain but results in the wrong infinite-coupling limit.

Overall, the pxLDA provides fairly accurate predictions for ultrastrong light-matter coupling with minimal additional costs when compared to common density-functional approximations for the Coulombic interaction. This computational simplicity allows us to utilize the pxLDA in an adiabatic manner also for time propagation. Fig. 6 presents the linear-response spectrum (similar to Fig. 2) and illustrates that the adiabatic pxLDA potential correctly predicts the upward-bending of the first matter excitation at low frequencies. However, it falsely predicts the same behavior for all excited states.

There are several possible origins for that. First, since in the time-dependent case we have complex wave functions, in the original Eq. 16 we should work with $\Delta \hat{\mathbf{J}}_{\text{p}}$ instead of $\hat{\mathbf{J}}_{\text{p}}$. Thus, Eq. 18 also includes the implicit approximation $\mathbf{J}_{\text{p}} \equiv 0$ due to the adiabatic homogeneous-electron-gas ansatz for the fluctuations. This can be corrected by working with Eq. 16 and by resubstituting $\Delta \hat{\mathbf{J}}_{\text{p}}$. Second, in the time-dependent case we should actually use a different equation of motion (total current instead of just the paramagnetic current), which leads to the contribution of several further terms that are zero in the static case, and we should use a consistent approximation to the Mx vector potential as well (SI Appendix). That is, to be consistent with the static approximation we should actually use a current-density formulation of QEDFT also in the dipole-coupling limit. In addition, we need to include nonadiabatic photon fluctuation effects. This poses a real problem for the simple Hamiltonian approach discussed in Section 1 as the substituted operators are defined in the Heisenberg picture which would lead to involved memory terms in order to

arrive at the photon-free Hamiltonian in the Schrödinger picture. Yet, for the equation-of-motion approach employed in this section, no such restriction applies as expectation values are by definition identical in all pictures, and we can easily switch between them. This allows us to define also a nonadiabatic version of the presented px potential (as discussed in *SI Appendix*) leading to multiphoton excitations.

Furthermore, for a detailed understanding of light-matter coupling effects, it is desirable to go beyond the common dipole approximation. Again we can simply follow the above strategy and replace the fluctuations of the vector-potential operator by inverting the inhomogeneous quantum Maxwell equation, which is discussed in *SI Appendix*. Yet since we can derive the defining equation for the px potential by taking the long-wavelength limit of the corresponding minimal-coupling equations (see also *SI Appendix*), we can find a first approximation to the beyond-dipole case by simply making the coupling terms $\omega_{d,\alpha} \propto \lambda_\alpha^2$ and the polarizations $\tilde{\epsilon}_\alpha$ spatially dependent in Eqs. 16 or 18. Such a simple extension would already lead to (static) ponderomotive forces that allow trapping of polarizable systems in the eigenmode profile. In order to practically apply our full minimal coupling extensions to extended systems we have to account efficiently for the new commensurable period of cavity mode and crystal structure, which will be in general substantially bigger.

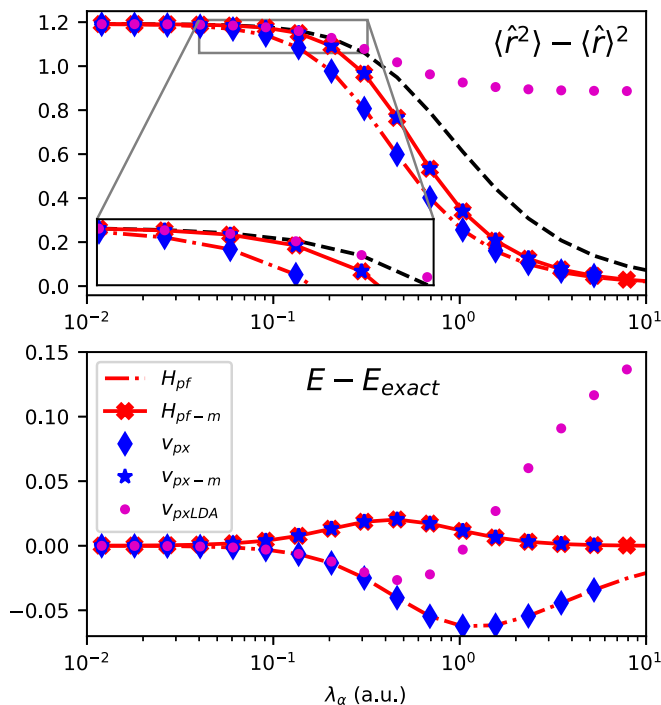


Fig. 5. (Top) Dipole variance $\langle \hat{r}^2 \rangle - \langle \hat{r} \rangle^2$ and (Bottom) total energy difference to the exact solution $E - E_{\text{exact}}$ of the correlated cavity hydrogen system for various approximations compared to the exact solution (black dashed line) with 40 photon number states (using the PZW Hamiltonian). The index “-m” indicates the utilization of the ad hoc mollification of the external potential according to Section 2. The mollification leads to variational energies. The exchange LDA potential provides excellent results up to $\lambda = 0.3$. The cavity frequency of the single cavity mode is set in resonance with the bare lowest excitation energy like in Fig. 2. The real-space grid and potential were chosen identically to Fig. 3. Since we selected particularly tough parameters for the benchmark, the approximations presented here will perform better for softer potentials (compare Section 2).

While supercell calculations provide a simple shortcut, they are computationally demanding such that adjusting the Bloch ansatz becomes particularly interesting (66).

Finally, we note that we have limited our initial investigation presented here to the exchange-level of theory in the longitudinal and the transverse light-matter interactions. Besides going beyond the single-Slater determinant ansatz (64, 67) or combinations with wave function-based methods (68), there is the possibility (hitherto only for the longitudinal interaction) to use directly parametrizations or approximations of correlation expressions (69, 70). The photon-free framework provides us with functionals that are in close structural relation to the known approximations and conditions developed in ordinary density-functional theory which facilitates the further development of QEDFT. Providing accurate yet computationally affordable functionals that go beyond the dipole approximation and include correlations and nonadiabatic contributions will be the subject of future work.

4. Conclusion

Solving the Schrödinger equation for realistic system is an extremely hard task. The moment we consider the quantum character of light, we add an abundance of additional degrees of freedom to the already almost intractable problem. Our goal to retain the first-principles character of electronic structure theory and entwine it with QED is thus particularly challenging, unless we find effective descriptions which allow us to absorb large parts of the QED problem into the familiar electronic problem.

We presented here such an effective photon-free framework which adheres to all essential physical demands, including the correct frequency and cavity polarization dependence and accounts even in its simplest form for a substantial part of the full QED problem. Even if we remain entirely within the electronic Hilbert space, we recover the exact results of light-matter interaction in the weak coupling, deep ultrastrong coupling, and high frequency limit, as well as for homogeneous systems. This framework provides an excellent electronic subspace solution which becomes exact whenever a factorized wave function ansatz is possible. Starting from there, we illustrated that the photon-free approach corresponds to a highly efficient basis for light-matter interaction in many relevant situations. Expressing the coupling purely in matter quantities provided a convenient starting point to develop exchange correlation potentials dealing with correlated light-matter systems. We then leveraged the potential of the photon-free ansatz by constructing a hierarchy of QEDFT functionals, illustrated their performance for ground and excited states, and discussed how to go beyond the simple adiabatic and dipole-approximated cases. As a result, we derived a local-density approximated functional that combines the quantum nature of light with the computational simplicity of well-known electronic local-density functionals in DFT. The photon-free construction, including the derived QEDFT functionals, is ideally suited to describe periodic systems strongly coupled to light.

That the photon-free QED Hamiltonian remains entirely within the electronic Hilbert space is of great benefit not only computationally but also conceptually. The latter feature allows it to serve as a foundation for the development of more advanced electron-photon-structure approaches without the need to consider extensions of the known methodologies to larger Fock spaces. For instance, the description of strong light-matter coupling in terms of nonequilibrium Green’s functions commonly introduces the necessity to solve an expanded set of Kadanoff-Baym equations (71). The present photon-free QED ansatz on the other hand could serve as a starting point that remains exact in the infinite coupling limit. It is thus ideal for perturbative improvements on top of its nonperturbative foundation.

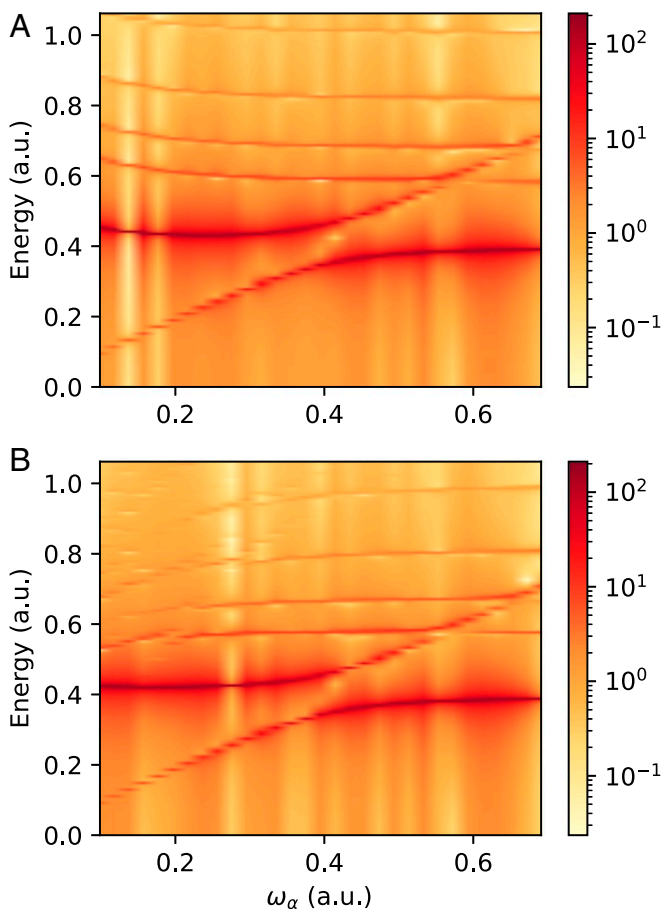


Fig. 6. Linear response spectrum of the dipole moment $|r(\omega_\alpha)|$ of a single cavity mode with varying frequency ω_α coupled to one-dimensional soft-Coulomb hydrogen. Shown are the adiabatic utilization of the pLDA potential Eq. 18 in combination with (A) the semiclassical Maxwell-Schrödinger solution and (B) the exact reference solution. All parameters are identical to Fig. 2.

Our approach is conceptually general enough to be expanded to vibrational light-matter coupling and beyond the dipole approximation. Extensions beyond the dipole approximation become feasible if we follow the equation-of-motion constructions of Section 3. The obtained spectral information and the lack of multiphoton excitations suggest that future development should foremost focus on going beyond the adiabatic approximation. While this demands curing an old wound of density-functional theory, the equation-of-motion construction provides a passage to memory-dependent functionals. It furthermore avoids the numerical and perturbative instability of other orbital-dependent functionals such as the time-dependent optimized-effective potential (72). This would not only greatly benefit the description of strongly correlated light-matter systems but also ameliorate common problems with ordinary density-functional theory.

5. Materials and Methods

All calculations have been performed using python3; the code can be obtained from the authors upon reasonable request.

5.1. The Bogoliubov Transformation. In the following, we recall the well-known Bogoliubov transformation (ref. 73, Section 1.10) utilized in Section 1. The transformation is based on the realization that the purely photonic part of Hamiltonian Eq. 1,

$$\hat{H}_{\text{ph}} = \sum_{\alpha=1}^{M_p} \omega_\alpha \left(\hat{a}_\alpha^\dagger \hat{a}_\alpha + \frac{1}{2} \right) + \frac{N_e}{2c^2} \hat{\mathbf{A}}^2,$$

with $\hat{\mathbf{A}}$ given by Eq. 2 is simply the Hamiltonian of M_p coupled harmonic oscillators. This is even more obvious when we introduce the harmonic oscillator coordinates $q_\alpha = 1/\sqrt{2\omega_\alpha} (\hat{a}_\alpha^\dagger + \hat{a}_\alpha)$ and $p_\alpha = i\sqrt{\omega_\alpha/2} (\hat{a}_\alpha^\dagger - \hat{a}_\alpha)$, which leads to the Hamiltonian

$$\hat{H}_{\text{ph}} = \frac{1}{2} \left(\sum_{\alpha=1}^{M_p} p_\alpha^2 + \sum_{\alpha, \alpha'=1}^{M_p} W_{\alpha, \alpha'} q_\alpha q_{\alpha'} \right),$$

with $W_{\alpha, \alpha'} = \omega_\alpha^2 \delta_{\alpha, \alpha'} + 4\pi N_e S_\alpha(\mathbf{r}_0) S_{\alpha'}(\mathbf{r}_0) \epsilon_\alpha \cdot \epsilon_{\alpha'}$. Since $W_{\alpha, \alpha'}$ is symmetric, there is a unitary transformation U which brings $W_{\alpha, \alpha'}$ into diagonal form $\tilde{\Omega} = U W U^\dagger$ with eigenvalues $\tilde{\omega}_\alpha^2$. This diagonalization introduces the decoupled normal modes with corresponding operators $\tilde{q}_\beta = \sum_\alpha U_{\beta, \alpha} q_\alpha$ and $\tilde{p}_\beta = \sum_\alpha U_{\beta, \alpha} p_\alpha$ and polarization vectors $\tilde{\epsilon}_\beta = \sum_\alpha U_{\beta, \alpha} \epsilon_\alpha$. The resulting Hamiltonian reads

$$\begin{aligned} \hat{H}_{\text{PF}} = & -\frac{1}{2} \sum_{i=1}^{N_e} \nabla_i^2 + \sum_{i=1}^{N_e} v(\mathbf{r}_i) + \frac{1}{2} \sum_{i \neq j}^{N_e} w(\mathbf{r}_i, \mathbf{r}_j) \\ & + \frac{1}{c} \hat{\mathbf{j}}_p \cdot \hat{\mathbf{A}} + \sum_{\alpha=1}^{M_p} \tilde{\omega}_\alpha \left(\hat{a}_\alpha^\dagger \hat{a}_\alpha + \frac{1}{2} \right), \end{aligned}$$

where we introduced the annihilation operators $\hat{a}_\alpha = 1/\sqrt{2\tilde{\omega}_\alpha} (\tilde{\omega}_\alpha \tilde{q}_\alpha + i\tilde{p}_\alpha)$ corresponding to $\tilde{q}_\alpha, \tilde{p}_\alpha$, as well as the respective creation operators. We will ignore the $\tilde{\cdot}$ denotation of operators for the sake of brevity and readability.

5.2. Adiabatic Breit Approximation for the Photon Energy. A sensible effective photon-free Hamiltonian (e.g., Eq. 7) has to adhere to the same fundamental rules as the original Hamiltonian. Foremost, this includes the existence of bound eigenstates and translational invariance. To achieve this, we cannot just disregard all photonic degrees of freedom in an ad hoc fashion but instead have to find the matter term corresponding to $\sum_\alpha \tilde{\omega}_\alpha \hat{a}_\alpha^\dagger \hat{a}_\alpha$. From the equations of motion we get for the annihilation operators (the creation operators are always just the Hermitian conjugate),

$$\frac{d^2}{dt^2} \hat{a}_\alpha = -\tilde{\omega}_\alpha^2 \hat{a}_\alpha - \frac{\omega_{d, \alpha} \sqrt{\tilde{\omega}_\alpha}}{\sqrt{2N_e}} \tilde{\epsilon}_\alpha \cdot \left(\hat{\mathbf{j}}_p - \frac{i}{\tilde{\omega}_\alpha} \frac{d}{dt} \hat{\mathbf{j}}_p \right),$$

and we can identify the adiabatic approximation as

$$\hat{a}_\alpha \approx -\frac{\omega_{d, \alpha}}{\sqrt{2N_e \tilde{\omega}_\alpha^3}} \tilde{\epsilon}_\alpha \cdot \hat{\mathbf{j}}_p.$$

This leads to the critical contribution that renders the photon-free Hamiltonian bounded from below

$$\tilde{\omega}_\alpha \hat{a}_\alpha^\dagger \hat{a}_\alpha \approx \frac{\omega_{d, \alpha}^2}{2N_e \tilde{\omega}_\alpha^2} (\tilde{\epsilon}_\alpha \cdot \hat{\mathbf{j}}_p)^2.$$

In the time-dependent case, the bilinear term $\hat{\mathbf{j}}_p \cdot \hat{\mathbf{A}} \rightarrow (\tilde{\epsilon}_\alpha \cdot \Delta \hat{\mathbf{j}}_p) (\tilde{\epsilon}_\alpha \cdot \hat{\mathbf{j}}_p)$ suggests that also the quadratic $\hat{a}_\alpha^\dagger \hat{a}_\alpha$ term should follow the same structure. This motivates the substitution $\tilde{\omega}_\alpha \hat{a}_\alpha^\dagger \hat{a}_\alpha \approx \omega_{d, \alpha}^2 / (2N_e \tilde{\omega}_\alpha^2) (\tilde{\epsilon}_\alpha \cdot \Delta \hat{\mathbf{j}}_p) (\tilde{\epsilon}_\alpha \cdot \hat{\mathbf{j}}_p)$ and leads to the photon-free Hamiltonian Eq. 7. Other possibilities, such as using the static (time-independent) substitution of $\hat{a}_\alpha^\dagger \hat{a}_\alpha$ or disregarding the mean-field correction in Eq. 7 $(\tilde{\epsilon}_\alpha \cdot \Delta \hat{\mathbf{j}}_p) (\tilde{\epsilon}_\alpha \cdot \hat{\mathbf{j}}_p) \rightarrow (\tilde{\epsilon}_\alpha \cdot \hat{\mathbf{j}}_p)^2$, have been investigated but provided worse results.

5.3. Photon Exchange-Only LDA Derivation. In order to arrive at a local-density approximation for the photon exchange-only term, we note that for a properly antisymmetrized wave function $\Phi(\mathbf{r}, \bar{\mathbf{r}})$, $\bar{\mathbf{r}} = (\mathbf{r}_2 \dots \mathbf{r}_{N_e})$, we can express the expectation value in Eq. 16 in terms of the one-body (1RDM) and two-body reduced density matrices (2RDM) (ref. 69, Section 2.4) as

$$\begin{aligned}
\mathbf{f}_\alpha(\mathbf{r}) &= \left\langle (\bar{\epsilon}_\alpha \cdot \hat{\mathbf{j}}_p) \hat{\mathbf{j}}_p(\mathbf{r}) \right\rangle = \left\langle (\bar{\epsilon}_\alpha \cdot \hat{\mathbf{j}}_p) \Phi, \hat{\mathbf{j}}_p(\mathbf{r}) \Phi \right\rangle \\
&= \frac{N_e}{2i} \int \left[\left((\bar{\epsilon}_\alpha \cdot \hat{\mathbf{j}}_p) \Phi \right)^* \nabla \Phi - \left((\bar{\epsilon}_\alpha \cdot \hat{\mathbf{j}}_p) \nabla \Phi \right)^* \Phi \right] d\mathbf{r} \\
&= \frac{N_e}{2} \int \left[\left((\bar{\epsilon}_\alpha \cdot \nabla) \Phi^* \right) \nabla \Phi - \left((\bar{\epsilon}_\alpha \cdot \nabla) \nabla \Phi^* \right) \Phi \right] d\mathbf{r} \\
&+ \frac{N_e(N_e - 1)}{2} \int \underbrace{\left[\left((\bar{\epsilon}_\alpha \cdot \nabla_2) \Phi^* \right) \nabla \Phi - \left((\bar{\epsilon}_\alpha \cdot \nabla_2) \nabla \Phi^* \right) \Phi \right]}_{((\bar{\epsilon}_\alpha \cdot \nabla_2) \Phi^*) \nabla \Phi + c.c.} d\mathbf{r} \quad [19] \\
&= \frac{1}{2} \left[(\bar{\epsilon}_\alpha \cdot \nabla') \nabla \rho_{(1)}(\mathbf{r}, \mathbf{r}') - (\bar{\epsilon}_\alpha \cdot \nabla') \nabla' \rho_{(1)}(\mathbf{r}, \mathbf{r}') \right]_{r'=r} \\
&+ \int \left[(\bar{\epsilon}_\alpha \cdot \nabla'_2) \nabla \rho_{(2)}(\mathbf{r}, \mathbf{r}_2; \mathbf{r}', \mathbf{r}'_2) + c.c. \right]_{r'=r, r'_2=r_2} d\mathbf{r}_2.
\end{aligned}$$

We used partial integration to move $\bar{\epsilon}_\alpha \cdot \nabla_2$ to the other side in the under-braced expression. This 2RDM contribution will later be found having the same form as the 1RDM terms but with opposite sign, lessening the strength of the attractive px potential, and will finally be neglected altogether. In the remaining integral the 2RDM can be rewritten in terms of the 1RDM for closed-shell Slater-determinant states of spin- $\frac{1}{2}$ particles that we assume from here, $\rho_{(2)}(\mathbf{r}, \mathbf{r}_2; \mathbf{r}', \mathbf{r}'_2) = \frac{1}{2} [\rho_{(1)}(\mathbf{r}, \mathbf{r}') \rho_{(1)}(\mathbf{r}_2, \mathbf{r}'_2) - \frac{1}{2} \rho_{(1)}(\mathbf{r}, \mathbf{r}'_2) \rho_{(1)}(\mathbf{r}_2, \mathbf{r}')]$. This makes the integral

$$\begin{aligned}
&\int (\bar{\epsilon}_\alpha \cdot \nabla'_2) \nabla \rho_{(2)}(\mathbf{r}, \mathbf{r}_2; \mathbf{r}', \mathbf{r}'_2) \Big|_{r'=r, r'_2=r_2} d\mathbf{r}_2 \\
&= \frac{1}{2} \int \left[\nabla \rho_{(1)}(\mathbf{r}, \mathbf{r}') \Big|_{r'=r} (\bar{\epsilon}_\alpha \cdot \nabla'_2) \rho_{(1)}(\mathbf{r}_2, \mathbf{r}'_2) \Big|_{r'_2=r_2} \right. \\
&\quad \left. - \frac{1}{2} \left((\bar{\epsilon}_\alpha \cdot \nabla_2) \nabla \rho_{(1)}(\mathbf{r}, \mathbf{r}_2) \right) \rho_{(1)}(\mathbf{r}_2, \mathbf{r}) \right] d\mathbf{r}_2 \quad [20] \\
&= \frac{1}{2} \nabla \rho_{(1)}(\mathbf{r}, \mathbf{r}') \Big|_{r'=r} \int (\bar{\epsilon}_\alpha \cdot \nabla'_2) \rho_{(1)}(\mathbf{r}_2, \mathbf{r}'_2) \Big|_{r'_2=r_2} d\mathbf{r}_2 \\
&\quad - \frac{1}{4} \int \left((\bar{\epsilon}_\alpha \cdot \nabla_2) \nabla \rho_{(1)}(\mathbf{r}, \mathbf{r}_2) \right) \rho_{(1)}(\mathbf{r}_2, \mathbf{r}) d\mathbf{r}_2.
\end{aligned}$$

For the homogeneous electron gas of spin- $\frac{1}{2}$ particles the 1RDM is then given by the usual Fermi sphere integration (ref. 69, Section 6.1)

$$\rho_{(1)}(\mathbf{r}, \mathbf{r}') = \frac{2}{(2\pi)^d} \int_{|\mathbf{k}| < k_F} e^{i\mathbf{k} \cdot (\mathbf{r} - \mathbf{r}')} d\mathbf{k}, \quad [21]$$

with the local Fermi radius given by $k_F(\mathbf{r}) = 2\pi(\rho(\mathbf{r})/(2V_d))^{1/d}$ with V_d being the volume of the d -dimensional unit sphere (see ref. 74, equation 1.77, but also easily derived from Eq. 21 by setting $\mathbf{r}' = \mathbf{r}$). The Fermi sphere is centered around the origin since we consider the static case with zero current $\hat{\mathbf{j}}_p = 0$. Then

$$\nabla \rho_{(1)}(\mathbf{r}, \mathbf{r}') \Big|_{r'=r} = \frac{2i}{(2\pi)^d} \int_{|\mathbf{k}| < k_F} \mathbf{k} d\mathbf{k} = 0$$

is readily seen to be zero because \mathbf{k} is integrated over the symmetric Fermi sphere volume. We now insert the ansatz Eq. 21 also into all remaining terms from Eq. 19 and get with $k_F = k_F(\mathbf{r})$, chosen at \mathbf{r} because this is the primary position, and k'_F left open that

$$\begin{aligned}
\mathbf{f}_\alpha(\mathbf{r}) &= \frac{2}{(2\pi)^d} \int_{|\mathbf{k}| < k_F} (\bar{\epsilon}_\alpha \cdot \mathbf{k}) \mathbf{k} d\mathbf{k} \\
&- \frac{1}{(2\pi)^{2d}} \iint_{\substack{|\mathbf{k}| < k_F \\ |\mathbf{k}'| < k'_F}} (\bar{\epsilon}_\alpha \cdot \mathbf{k}) \mathbf{k} \left(\int e^{i\mathbf{k} \cdot (\mathbf{r} - \mathbf{r}_2)} e^{i\mathbf{k}' \cdot (\mathbf{r}_2 - \mathbf{r})} d\mathbf{r}_2 + c.c. \right) d\mathbf{k} d\mathbf{k}' \\
&= \frac{2}{(2\pi)^d} \int_{|\mathbf{k}| < k_F} (\bar{\epsilon}_\alpha \cdot \mathbf{k}) \mathbf{k} d\mathbf{k} - \frac{2}{(2\pi)^d} \int_{|\mathbf{k}| < \min(k_F, k'_F)} (\bar{\epsilon}_\alpha \cdot \mathbf{k}) \mathbf{k} d\mathbf{k}.
\end{aligned}$$

Here the \mathbf{r}_2 integration leads to a delta function that leaves the smaller integration radius from the two \mathbf{k}, \mathbf{k}' integrals. What is done here is to introduce the local approximation Eq. 21 for $\rho_{(1)}(\mathbf{r}, \mathbf{r}_2)$ even though the positions \mathbf{r}, \mathbf{r}_2 are not close. Since further \mathbf{r}_2 is taken from the whole space, we can argue that for an inhomogeneous medium, $\min(k_F, k'_F)$ approaches zero because k'_F possibly gets very small, and thus, no contribution arises from the original 2RDM expression. On the other hand, for a homogeneous medium it will hold $\min(k_F, k'_F) = k_F$ since $k'_F = k_F$, and thus, $\mathbf{f}_\alpha(\mathbf{r}) = 0$ which exactly fits

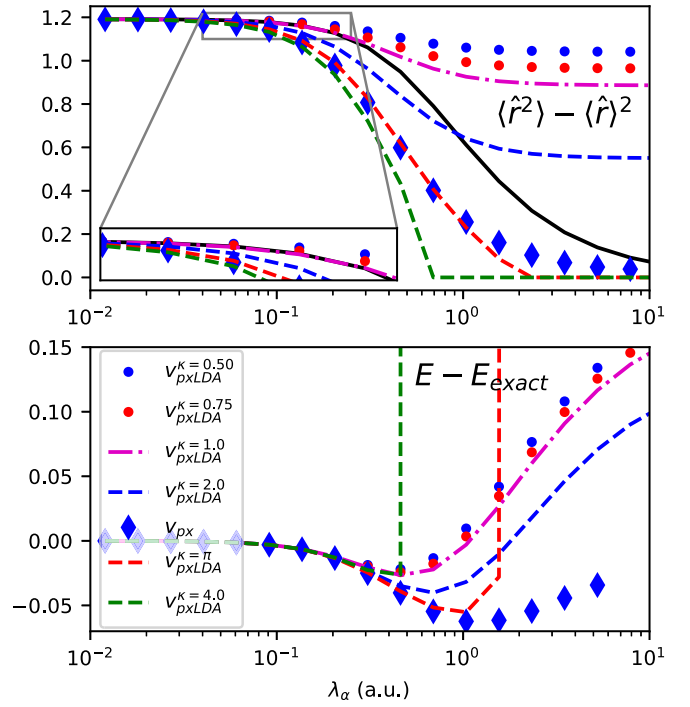


Fig. 7. (Top) Dipole variance and (Bottom) total energy difference to the exact solution for the exchange LDA approximation V_{pxLDA}^{κ} with different parameters κ compared to the exact reference solution (black solid) for the one-dimensional soft-Coulomb potential with $\xi = 1$ coupled to a single cavity mode in resonance to the first bare excitation energy. The setup is identical to Fig. 5. The choice $\kappa = 1$ suggested here provides excellent results up to $\lambda = 0.3$. Larger values can improve the (deep) ultrastrong coupling limit at the cost of instability (collapse onto a few/single grid point) and overestimation in the strong to ultrastrong domain.

our expectations. To take all those different situations into account, we introduce a factor $\kappa \in [0, 1]$ that expresses the ratio that is left from the first integral after subtracting the second with a smaller radius. The maximally inhomogeneous limit corresponds to $\kappa = 1$ while the homogeneous case is $\kappa = 0$. Within the main text we decided to limit ourselves entirely to the case $\kappa = 1$. Considering spin-polarized systems motivates a spin-resolved LDA which would suggest a different κ regime and will be the subject of future work. Introducing the new parameter κ is a simple approach that allows us to stay in the realm of the local density approximation. Clearly, those considerations suggest that the light-matter interaction, as it is non-local in character, should be described ideally by more advanced nonlocal functionals following the spirit of modern (meta) GGAs. Fig. 7 illustrates the performance of the LDA for various values also including the possibility $\kappa > 1$. In this example, the LDA potential with $\kappa = 1$ provides even better results than the full exchange potential for $\lambda < 0.3$ due to error compensation.

Evaluating the integral by switching to polar coordinates we get

$$\mathbf{f}_\alpha(\mathbf{r}) = \frac{2\kappa}{(2\pi)^d} \int_{|\mathbf{k}| < k_F} (\bar{\epsilon}_\alpha \cdot \mathbf{k}) \mathbf{k} d\mathbf{k} = \frac{2\kappa V_d k_F(\mathbf{r})^{d+2}}{(2\pi)^d} \frac{d+2}{d+2} \bar{\epsilon}_\alpha.$$

Putting this solution for $\mathbf{f}_\alpha(\mathbf{r})$ into Eq. 16 yields a Poisson-like equation for the photon exchange-only LDA potential,

$$\begin{aligned}
\nabla^2 V_{pxLDA}(\mathbf{r}) &= -\nabla \cdot \left[\sum_{\alpha=1}^{M_p} \frac{\omega_{d,\alpha}^2}{N_e \omega_\alpha^2} (\bar{\epsilon}_\alpha \cdot \nabla) \mathbf{f}_\alpha(\mathbf{r}) \right] \\
&= -\frac{2\kappa V_d}{(2\pi)^d} \sum_{\alpha=1}^{M_p} \frac{\omega_{d,\alpha}^2}{N_e \omega_\alpha^2} (\bar{\epsilon}_\alpha \cdot \nabla) \left[\frac{k_F(\mathbf{r})^{d+1}}{\rho(\mathbf{r})} (\bar{\epsilon}_\alpha \cdot \nabla) k_F(\mathbf{r}) \right].
\end{aligned}$$

Expressing the Fermi radius by the local density once more, we get the final result

$$\nabla^2 V_{\text{pxLDA}}(\mathbf{r}) = - \sum_{\alpha=1}^{M_p} \frac{2\kappa\pi^2\omega_{d,\alpha}^2}{N_e\bar{\omega}_{\alpha}^2} (\bar{\epsilon}_{\alpha} \cdot \nabla)^2 \left(\frac{\rho(\mathbf{r})}{2V_d} \right)^{\frac{2}{d}}$$

Assuming an isotropic mode distribution, the differential operators $(\bar{\epsilon}_{\alpha} \cdot \nabla)^2$ are summed up to full Laplacians, and thus, if zero boundary conditions are assumed, the photon exchange-only LDA potential is explicitly given by

$$V_{\text{pxLDA}}(\mathbf{r}) = - \sum_{\alpha=1}^{M_p} \frac{2\kappa\pi^2\omega_{d,\alpha}^2}{dN_e\bar{\omega}_{\alpha}^2} \left(\frac{\rho(\mathbf{r})}{2V_d} \right)^{\frac{2}{d}}$$

Data Availability. All study data are included in the article and *SI Appendix*.

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1. T. W. Ebbesen, Hybrid light-matter states in a molecular and material science perspective. *Acc. Chem. Res.* **49**, 2403–2412 (2016).
2. R. Chikkaraddy *et al.*, Single-molecule strong coupling at room temperature in plasmonic nanocavities. *Nature* **535**, 127–130 (2016).
3. A. F. Kockum, A. Miranowicz, S. De Liberato, S. Savasta, F. Nori, Ultrastrong coupling between light and matter. *Nat. Rev. Phys.* **1**, 19 (2019).
4. D. N. Basov, A. Asenjo-Garcia, P. J. Schuck, X. Zhu, A. Rubio, Polariton panorama. *Nanophotonics* **10**, 549–577 (2021).
5. J. A. Hutchison, T. Schwartz, C. Genet, E. Devaux, T. W. Ebbesen, Modifying chemical landscapes by coupling to vacuum fields. *Angew. Chem. Int. Ed.* **51**, 1592–1596 (2012).
6. A. Thomas *et al.*, Ground-state chemical reactivity under vibrational coupling to the vacuum electromagnetic field. *Angew. Chem. Int. Ed. Engl.* **55**, 11462–11466 (2016).
7. B. Munkhbat, M. Wersäll, D. G. Baranov, T. J. Antosiewicz, T. Shegai, Suppression of photo-oxidation of organic chromophores by strong coupling to plasmonic nanoantennas. *Sci. Adv.* **4**, eaas9552 (2018).
8. J. Galego, F. J. Garcia-Vidal, J. Feist, Cavity-induced modifications of molecular structure in the strong-coupling regime. *Phys. Rev. X* **5**, 041022 (2015).
9. F. Herrera, F. C. Spano, Cavity-controlled chemistry in molecular ensembles. *Phys. Rev. Lett.* **116**, 238301 (2016).
10. G. Groenhof, C. Climent, J. Feist, D. Morozov, J. J. Toppari, Tracking polariton relaxation with multiscale molecular dynamics simulations. *J. Phys. Chem. Lett.* **10**, 5476–5483 (2019).
11. C. Schäfer, J. Flick, E. Ronca, P. Narang, A. Rubio, Shining light on the microscopic resonant mechanism responsible for cavity-mediated chemical reactivity. *arXiv [Preprint]* (2021). <https://arxiv.org/abs/2104.12429> (Accessed 18 June 2021).
12. D. M. Coles *et al.*, Polariton-mediated energy transfer between organic dyes in a strongly coupled optical microcavity. *Nat. Mater.* **13**, 712–719 (2014).
13. E. Orghiu *et al.*, Conductivity in organic semiconductors hybridized with the vacuum field. *Nat. Mater.* **14**, 1123–1129 (2015).
14. X. Zhong *et al.*, Energy transfer between spatially separated entangled molecules. *Angew. Chem. Int. Ed. Engl.* **56**, 9034–9038 (2017).
15. J. Schachenmayer, C. Genes, E. Tignone, G. Pupillo, Cavity-enhanced transport of excitons. *Phys. Rev. Lett.* **114**, 196403 (2015).
16. J. Feist, F. J. Garcia-Vidal, Extraordinary exciton conductance induced by strong coupling. *Phys. Rev. Lett.* **114**, 196402 (2015).
17. R. Sáez-Blázquez, J. Feist, A. I. Fernández-Domínguez, F. J. García-Vidal, Organic polaritons enable local vibrations to drive long-range energy transfer. *Phys. Rev. B* **97**, 241407 (2018).
18. M. Du *et al.*, Theory for polariton-assisted remote energy transfer. *Chem. Sci. (Camb.)* **9**, 6659–6669 (2018).
19. C. Schäfer, M. Ruggenthaler, H. Appel, A. Rubio, Modification of excitation and charge transfer in cavity quantum-electrodynamical chemistry. *Proc. Natl. Acad. Sci. U.S.A.* **116**, 4883–4892 (2019).
20. A. Thomas, *et al.*, Exploring superconductivity under strong coupling with the vacuum electromagnetic field. *arXiv [Preprint]* (2019). <https://arxiv.org/abs/1911.01459> (Accessed 25 May 2020).
21. M. A. Sentef, M. Ruggenthaler, A. Rubio, Cavity quantum-electrodynamical polaritonically enhanced electron-phonon coupling and its influence on superconductivity. *Sci. Adv.* **4**, eaau6969 (2018).
22. F. Schlawin, A. Cavalleri, D. Jaksch, Cavity-mediated electron-photon superconductivity. *Phys. Rev. Lett.* **122**, 133602 (2019).
23. D. G. Baranov, M. Wersäll, J. Cuadra, T. J. Antosiewicz, T. Shegai, Novel nanostructures and materials for strong light-matter interactions. *ACS Photonics* **5**, 24–42 (2018).
24. P. Forn-Díaz, L. Lamata, E. Rico, J. Kono, E. Solano, Ultrastrong coupling regimes of light-matter interaction. *Rev. Mod. Phys.* **91**, 025005 (2019).
25. N. S. Mueller *et al.*, Deep strong light-matter coupling in plasmonic nanoparticle crystals. *Nature* **583**, 780–784 (2020).
26. M. Hertzog, B. Munkhbat, D. Baranov, T. Shegai, K. Börjesson, Enhancing vibrational light-matter coupling strength beyond the molecular concentration limit using plasmonic arrays. *Nano Lett.* **21**, 1320–1326 (2021).
27. A. Goban *et al.*, Superradiance for atoms trapped along a photonic crystal waveguide. *Phys. Rev. Lett.* **115**, 063601 (2015).
28. N. K. Langford *et al.*, Experimentally simulating the dynamics of quantum light and matter at deep-strong coupling. *Nat. Commun.* **8**, 1715 (2017).
29. P. Türschmann *et al.*, Coherent nonlinear optics of quantum emitters in nanophotonic waveguides. *Nanophotonics* **8**, 1641–1657 (2019).
30. J. A. Campos-Gonzalez-Angulo, J. Yuen-Zhou, Polaritonic normal modes in transition state theory. *J. Chem. Phys.* **152**, 161101 (2020).
31. C. Climent, J. Feist, On the S_N2 reactions modified in vibrational strong coupling experiments: Reaction mechanisms and vibrational mode assignments. *Phys. Chem. Chem. Phys.* **22**, 23545–23552 (2020).
32. X. Li, A. Mandal, P. Huo, Cavity frequency-dependent theory for vibrational polariton chemistry. *Nat. Commun.* **12**, 1315 (2021).
33. C. Cohen-Tannoudji, J. Dupont-Roc, G. Grynberg, *Photons and Atoms—Introduction to Quantum Electrodynamics* (Wiley-VCH, 1997).
34. B. M. Garraway, The Dicke model in quantum optics: Dicke model revisited. *Philos. Trans. Royal Soc., Math. Phys. Eng. Sci.* **369**, 1137–1155 (2011).
35. D. Craig, T. Thirunamachandran, *Molecular Quantum Electrodynamics: An Introduction to Radiation-Molecule Interactions*, Dover Books on Chemistry Series (Dover Publications, 1998).
36. T. S. Haugland, C. Schäfer, E. Ronca, A. Rubio, H. Koch, Intermolecular interactions in optical cavities: An ab initio QED study. *J. Chem. Phys.* **154**, 094113 (2021).
37. M. Ruggenthaler *et al.*, Quantum-electrodynamical density-functional theory: Bridging quantum optics and electronic-structure theory. *Phys. Rev. A* **90**, 012508 (2014).
38. I. V. Tokatly, Time-dependent density functional theory for many-electron systems interacting with cavity photons. *Phys. Rev. Lett.* **110**, 233001 (2013).
39. M. Ruggenthaler, Ground-state quantum-electrodynamical density-functional theory. *arXiv [Preprint]* (2015). <https://arxiv.org/abs/1509.01417> (Accessed 25 May 2020).
40. R. Jestädt, M. Ruggenthaler, M. J. Oliveira, A. Rubio, H. Appel, Light-matter interactions within the Ehrenfest–Maxwell–Pauli–Kohn–Sham framework: Fundamentals, implementation, and nano-optical applications. *Adv. Phys.* **68**, 225–333 (2019).
41. C. Pellegrini, J. Flick, I. V. Tokatly, H. Appel, A. Rubio, Optimized effective potential for quantum electrodynamic time-dependent density functional theory. *Phys. Rev. Lett.* **115**, 093001 (2015).
42. J. Flick, C. Schäfer, M. Ruggenthaler, H. Appel, A. Rubio, Ab initio optimized effective potentials for real molecules in optical cavities: Photon contributions to the molecular ground state. *ACS Photonics* **5**, 992–1005 (2018).
43. J. H. Shirley, Solution of the Schrödinger equation with a Hamiltonian periodic in time. *Phys. Rev.* **138**, B979 (1965).
44. C. Schäfer, M. Ruggenthaler, A. Rubio, Ab initio nonrelativistic quantum electrodynamics: Bridging quantum chemistry and quantum optics from weak to strong coupling. *Phys. Rev. A (Coll. Park)* **98**, 043801 (2018).
45. U. De Giovannini, H. Hübener, Floquet analysis of excitations in materials. *J. Phys. Mater.* **3**, 012001 (2019).
46. G. Breit, Dirac’s equation and the spin-spin interactions of two electrons. *Phys. Rev.* **39**, 616–624 (1932).
47. V. Rokaj, M. Ruggenthaler, F. G. Eich, A. Rubio, The free electron gas in cavity quantum electrodynamics. *arXiv [Preprint]* (2020). <https://arxiv.org/abs/2006.09236> (Accessed 25 May 2021).
48. M. Griesemer, C. Tix, Instability of a pseudo-relativistic model of matter with self-generated magnetic field. *J. Math. Phys.* **40**, 1780–1791 (1999).
49. E. A. Power, S. Zienau, Coulomb gauge in non-relativistic quantum electro-dynamics and the shape of spectral lines. *Philos. Trans. Royal Soc. A* **251**, 427–454 (1959).
50. D. L. Andrews, G. A. Jones, A. Salam, R. G. Woolley, Perspective: Quantum Hamiltonians for optical interactions. *J. Chem. Phys.* **148**, 040901 (2018).
51. C. Schäfer, M. Ruggenthaler, V. Rokaj, A. Rubio, Relevance of the quadratic diamagnetic and self-polarization terms in cavity quantum electrodynamics. *ACS Photonics* **7**, 975–990 (2020).
52. V. Rokaj, D. M. Welakuh, M. Ruggenthaler, A. Rubio, Light-matter interaction in the long-wavelength limit: No ground-state without dipole self-energy. *J. Phys. B* **51**, 034005 (2018).
53. T. S. Haugland, E. Ronca, E. F. Kjønstad, A. Rubio, H. Koch, Coupled cluster theory for molecular polaritons: Changing ground and excited states. *Phys. Rev. X* **10**, 041043 (2020).
54. K. E. Cahill, R. J. Glauber, Ordered expansions in boson amplitude operators. *Phys. Rev.* **177**, 1857–1881 (1969).
55. Y. Ashida, A. İmamoğlu, E. Demler, Cavity quantum electrodynamics at arbitrary light-matter coupling strengths. *Phys. Rev. Lett.* **126**, 153603 (2021).
56. M. Ruggenthaler, F. Mackenroth, D. Bauer, Time-dependent Kohn-Sham approach to quantum electrodynamics. *Phys. Rev. A* **84**, 042107 (2011).

57. A. Pribram-Jones, D. A. Gross, K. Burke, DFT: A theory full of holes? *Annu. Rev. Phys. Chem.* **66**, 283–304 (2015).
58. M. M. Tchenkoue, M. Penz, I. Theophilou, M. Ruggenthaler, A. Rubio, Force balance approach for advanced approximations in density functional theories. *J. Chem. Phys.* **151**, 154107 (2019).
59. J. Flick, Simple exchange-correlation energy functionals for strongly coupled light-matter systems based on the fluctuation-dissipation theorem. *arXiv [Preprint]* (2021). <https://arxiv.org/abs/2104.06980> (Accessed 25 May 2021).
60. I. V. Tokatly, Quantum many-body dynamics in a Lagrangian frame: I. Equations of motion and conservation laws. *Phys. Rev. B Condens. Matter Mater. Phys.* **71**, 165104 (2005).
61. I. V. Tokatly, Quantum many-body dynamics in a Lagrangian frame: II. Geometric formulation of time-dependent density functional theory. *Phys. Rev. B Condens. Matter Mater. Phys.* **71**, 165105 (2005).
62. I. V. Tokatly, Time-dependent current density functional theory via time-dependent deformation functional theory: A constrained search formulation in the time domain. *Phys. Chem. Chem. Phys.* **11**, 4621–4630 (2009).
63. G. Vignale, C. A. Ullrich, S. Conti, Time-dependent density functional theory beyond the adiabatic local density approximation. *Phys. Rev. Lett.* **79**, 4878–4881 (1997).
64. M. Ruggenthaler, D. Bauer, Local Hartree-exchange and correlation potential defined by local force equations. *Phys. Rev. A* **80**, 052502 (2009).
65. N. Tancogne-Dejean *et al.*, Octopus, a computational framework for exploring light-driven phenomena and quantum dynamics in extended and finite systems. *J. Chem. Phys.* **152**, 124119 (2020).
66. T. Müller, S. Sharma, E. K. U. Gross, J. K. Dewhurst, Extending solid-state calculations to ultra-long-range length scales. *Phys. Rev. Lett.* **125**, 256402 (2020).
67. J. I. Fuks, S. E. B. Nielsen, M. Ruggenthaler, N. T. Maitra, Time-dependent density functional theory beyond Kohn-Sham Slater determinants. *Phys. Chem. Chem. Phys.* **18**, 20976–20985 (2016).
68. U. Mordovina, T. E. Reinhard, I. Theophilou, H. Appel, A. Rubio, Self-consistent density-functional embedding: A novel approach for density-functional approximations. *J. Chem. Theory Comput.* **15**, 5209–5220 (2019).
69. R. G. Parr, W. Yang, *Density Functional Theory of Atoms and Molecules* (Oxford University Press, 1989).
70. K. J. H. Giesbertz, R. van Leeuwen, U. von Barth, Towards nonlocal density functionals by explicit modeling of the exchange-correlation hole in inhomogeneous systems. *Phys. Rev. A* **87**, 022514 (2013).
71. G. D. Mahan, *Many-Particle Physics* (Springer Science & Business Media, 2013).
72. H. O. Wijewardane, C. A. Ullrich, Real-time electron dynamics with exact-exchange time-dependent density-functional theory. *Phys. Rev. Lett.* **100**, 056404 (2008).
73. F. H. Faisal, *Theory of Multiphoton Processes* (Springer, Berlin, 1987).
74. G. Giuliani, G. Vignale, *Quantum Theory of the Electron Liquid* (Cambridge University Press, 2005).