

# Theory and modeling of light-matter interactions in chemistry: current and future<sup>†</sup>

Braden M. Weight<sup>a,b</sup>, Xinyang Li,<sup>a</sup>, and Yu Zhang<sup>\*‡a</sup>

Light-matter interaction not only plays an instrumental role in characterizing the material's properties via various spectroscopic techniques but also provides a general strategy to manipulate material properties via the design of novel nanostructures. This perspective summarizes recent theoretical advances in modeling light-matter interactions in chemistry, mainly focusing on plasmon chemistry and polariton chemistry. The former utilizes the highly localized photon, plasmonic hot electrons, and local heat to drive chemical reactions. In contrast, polariton chemistry tune chemistry by modifying the potential energy curvatures via polariton formation. The perspective starts with the basic background of light-matter interactions, molecular quantum electrodynamics theory, and the challenges of modeling light-matter interactions in chemistry. Then, the recent advances in modeling plasmon and polariton chemistry are described, and future directions toward multiscale simulations of light-matter interaction-mediated chemistry are discussed.

## 1 Introduction

In 2007, a report compiled for the Office of Science of the Department of Energy (DOE) identified five grand challenges in basic energy science<sup>1</sup>, including 1) controlling material processes at the level of electrons, 2) designing and perfecting atom- and energy-efficient syntheses of new forms of matter with tailored properties, 3) understanding and controlling the remarkable properties of matter that emerge from complex correlations of atomic and electronic constituents, 4) mastering energy and information on the nanoscale to create new technologies with capabilities rivaling those of living things, and 5) characterizing and controlling matter away—especially far away—from equilibrium. One of the emerging techniques to address (some of) these challenges is light-matter interaction, which can be used to monitor, manipulate, and design materials' properties through multiple interactions between electrons, photons, and phonons<sup>2,3</sup>.

In conventional chemistry, chemists modify the functionalities of molecules based on different functional groups. For centuries, such a vision has been widely used in chemical synthesis to modify the chemical and physical properties of molecules or molecular materials. However, more is different<sup>4</sup>, and chemistry in a complex electromagnetic (EM) environment has emergent properties due to multiple couplings that can be used to control chemistry. Light-matter interactions have been instrumental in many branches of physics, chemistry, materials, and energy science<sup>3,5,6</sup>. In most of the previous applications, light-matter interaction is within the weak coupling scenario and can usually be treated at the lowest order in quantum electrodynamics via many-body perturbation theory<sup>7</sup>. Such treatments are widely used in different types of spectroscopy techniques<sup>8,9</sup>, optoelectronics<sup>10–14</sup>, quantum sensing<sup>15</sup>, quantum information<sup>16</sup>, light harvesting<sup>17,18</sup>, and beyond.

Alternatively, light-matter interactions open multiple new avenues for manipulating matter through novel emerging elementary excitations<sup>19</sup>, including plasmons and polaritons. Plas-

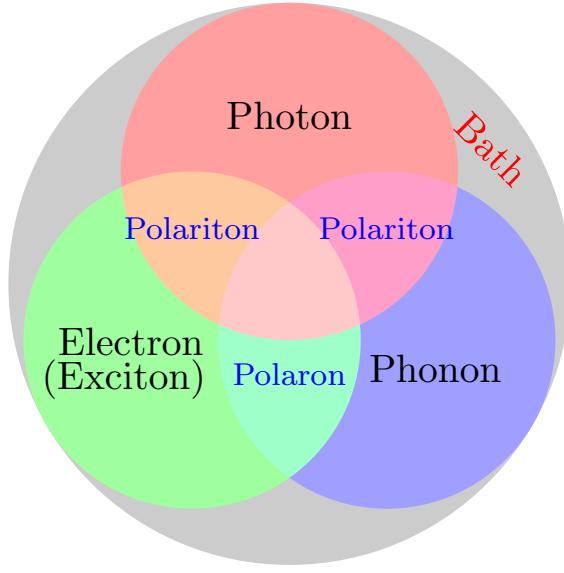
mons (either surface plasmons or localized surface plasmons) are the collective oscillations of conduction electrons in nanostructures, which can be excited when the frequency of external light matches the plasmon resonant energies. Plasmon excitation results in significantly amplified absorption and scattering cross-sections. Moreover, plasmon excitations can overcome the diffraction limit and concentrate the incident light into a highly localized volume, enhancing the electromagnetic (EM) field (or photons) by several orders of magnitude in the near field. The subsequent plasmon decay process generates hot electrons or heat through electron-electron and electron-phonon scatterings at different length and time scales<sup>20</sup>. Nevertheless, the resulting locally enhanced EM field, hot electrons, and heat can stimulate chemical reactions through various mechanisms<sup>21,22</sup>. In fact, plasmon-mediated chemical reactions (namely plasmon chemistry) have become a promising strategy to drive chemical processes over the past decade<sup>21</sup>.

On the other hand, when molecules are collectively and resonantly coupled with plasmon excitation, a new quasiparticle (namely polaritons) can be formed in the strong coupling regime. The term "strong" coupling is relative, meaning that the light-matter coupling is large enough to compete with or overcome dissipation or dephasing (i.e., when the coherent energy exchange between a confined light mode and quantum matter is faster than the decay and decoherence time scales of each part). Such strong coupling can be achieved when either a plasmonic mode is coupled with a few molecules<sup>23</sup> or many molecules are collectively coupled to a single cavity mode<sup>24</sup>. In the strong coupling regime, photons and electronic/excitonic excitations in matter become equally important and are strongly coupled on an equal quantized footing. As a consequence, individual "free" particles no longer exist. Instead, the fundamental excitations of the light-matter interacting system are polaritons, which are hybrid light-matter excitations (superpositions of quantized light and matter) (Fig 1a)<sup>3</sup> and possess both light and matter characteristics/topologies. Experiments have shown that matter properties can be modified with the formed polaritons, resulting in different photo-physics and photochemistry<sup>25</sup>. Since photon energies and light-matter coupling strengths are relatively tunable through cavity control, light-matter interaction in the strong coupling regime

<sup>a</sup> Theoretical Division, Los Alamos National Laboratory, Los Alamos, NM, 87545, USA.  
Tel: +1 505 606 2149; E-mail: zhy@lanl.gov

<sup>b</sup> Department of Physics and Astronomy, University of Rochester, Rochester, NY, 14627, U.S.A.

provides a fundamentally new way to manipulate matter properties for various desired applications, including lasing<sup>26,27</sup>, electronics<sup>28</sup>, long-range energy transfer<sup>29–32</sup>, Bose-Einstein condensates<sup>33–35</sup>, and chemical reactions<sup>36,37,37–56</sup>. Nevertheless, in polariton chemistry, light and matter cannot be treated as separate entities, as the strong coupling between them dresses each of them. Consequently, previous quantum chemistry methods for treating the electronic structure of matter become invalid. Brand new theoretical and modeling capabilities for describing polariton chemistry are required.



**Fig. 1** Light-matter interaction is a fundamentally multiscale/multiphysics problem. It involves multiple interactions among electrons, photons, and phonons at different time and length scales. As a result, multiple quasiparticles may be excited due to the light-matter interactions, including exciton (due to electron-hole interaction), (excitonic/vibrational) polariton (due to coupling between photon and exciton/phonon), polaron (due to electron-phonon coupling), and polaron-polaritons. In addition, all quantum systems are fundamentally open systems, leading to dissipation/dephasing due to the bath.

Despite the attractive applications of strong light-matter interaction, there are many open questions and fundamental problems about the mechanisms of physics and chemistry mediated by the strong light-matter interaction. The experimental progress should be complemented by theoretical advances. Progress in understanding the coupling between photons and elementary quasiparticles (plasmons, phonons, and excitons) in materials requires a generalized treatment of photons as one of the core degrees of freedom (DOFs) in light-matter interaction. Unfortunately, the strong light-matter interaction is fundamentally a multiscale and multiphysics problem that involves multiple interactions between many DOFs and their interplay with environments across different spatial and time scales. The traditional perturbation methods in the weak coupling regime are not applicable to the strong coupling regime, making it urgent to develop new theoretical and modeling techniques, especially multiscale methods, to understand and ultimately predict strong light-matter interaction-mediated physics and chemistry.

This perspective reviews recent theoretical efforts toward un-

derstanding the underlying mechanisms of plasmon and polariton chemistry due to the complex light-matter interactions and discusses our thoughts on future work. The article is structured as follows: Section 2 introduces the framework and basic mathematical structure of molecular quantum electrodynamics theory (QED), Section 3 explores the theoretical development in simulating plasmonic cavities and their strong interactions with molecules, Section 4 surveys recent progress in polariton chemistry in Fabry-Pérot-like cavities, and Section 5 concludes the discussion and provides a final perspective on future work in all realms of strong light-matter interaction.

## 2 Brief introduction to light-matter interactions and molecular QED theory

This section briefly introduces the quantum theories of light-matter interactions in the nonrelativistic limit<sup>57</sup>. All light-matter interactions arise from the interplay of matter DOFs (degrees of freedom) (electrons or spins, nuclei) and an EM environment. Hence, a full ab initio theory for light-matter interactions fundamentally requires the electronic structure theory of matter and principles of electrodynamics. Quantum electrodynamics (QED) is the indispensable and most precise theory for describing the interactions of charged particles and the dynamics of the EM field in mutual interaction.

### 2.1 Molecular quantum electrodynamics theory

The minimally coupled Coulomb Hamiltonian governs the non-relativistic dynamics of matter in an EM environment,

$$\hat{H} = \sum_i^{N_e+N_n} \frac{1}{2m_i} [\hat{P}_i - z_i \mathbf{A}(\mathbf{r}_i t)/c]^2 + \hat{V} + \hat{H}_{EM}, \quad (1)$$

with the Coulomb gauge  $\nabla \cdot \mathbf{A} = 0$ .  $\hat{P}_i = -i\hbar\nabla_i$  and  $z_i$  are the momentum operator and charge of particle  $i$ , respectively.  $N_e$  and  $N_n$  are the number of electrons and nuclei in the systems.  $\hat{V} = \hat{V}_{ee} + \hat{V}_{nn} + \hat{V}_{en} + V_{ext}$  contains all of the Coulomb interactions ( $\frac{1}{8\pi\epsilon_0} \frac{q_i q_j}{|\mathbf{r}_i - \mathbf{r}_j|}$ ) between the electronic and nuclear DOFs and another external potential. The Hamiltonian that describes the EM fields is

$$\hat{H}_{EM} = \frac{\epsilon_0}{2} \int d\mathbf{r} [\hat{\mathbf{E}}^2(\mathbf{r}) + c^2 \hat{\mathbf{B}}^2(\mathbf{r})]. \quad (2)$$

It obeys Maxwell's equations of motion and couples with the Schrödinger equation self-consistent through the vector potential  $\mathbf{A}(\mathbf{r}_i)$ . With multiple modes available, each field (including the vector potential  $\hat{\mathbf{A}}$ ) can be expressed as a sum over all possible radiation modes<sup>58,59</sup>

$$\hat{\mathbf{A}}(\mathbf{r}, t) = \sum_{\alpha} \mathbf{u}_{\alpha} (\hat{A}_{\alpha} e^{i(\mathbf{K} \cdot \mathbf{r} - \omega_{\alpha} t)} + \hat{A}_{\alpha}^{\dagger} e^{i(\mathbf{K} \cdot \mathbf{r} + \omega_{\alpha} t)}) \quad (3)$$

$$\hat{\mathbf{E}}(\mathbf{r}, t) = i \sum_{\alpha} \omega_{\alpha} \mathbf{u}_{\alpha} (\hat{A}_{\alpha} e^{i(\mathbf{K} \cdot \mathbf{r} - \omega_{\alpha} t)} + \hat{A}_{\alpha}^{\dagger} e^{i(\mathbf{K} \cdot \mathbf{r} + \omega_{\alpha} t)}) \quad (4)$$

$$\hat{\mathbf{B}}(\mathbf{r}, t) = i \sum_{\alpha} (\nabla \times \mathbf{u}_{\alpha}) (\hat{A}_{\alpha} e^{i(\mathbf{K} \cdot \mathbf{r} - \omega_{\alpha} t)} + \hat{A}_{\alpha}^{\dagger} e^{i(\mathbf{K} \cdot \mathbf{r} + \omega_{\alpha} t)}) \quad (5)$$

where  $\nabla \times \mathbf{u} = \mathbf{K} \times \mathbf{u}$ ,  $\alpha \equiv \lambda \mathbf{K}$  denotes the photon mode with momentum  $\mathbf{K}$  and polarization  $\lambda \in \{-1, 1\}$ ,  $\mathbf{u}$  is the unit vector de-

noting the direction of the vector potential, and  $\hat{A}_\alpha$  ( $\hat{A}_\alpha^\dagger$ ) is the mode decomposition coefficients that create (annihilate) the  $\alpha^{\text{th}}$  radiation mode. Note that  $\hat{\mathbf{E}} = -\frac{1}{c}\partial_t \hat{\mathbf{A}}(\mathbf{r})$  and  $\hat{\mathbf{B}} = \frac{1}{c}\nabla \times \hat{\mathbf{A}}(\mathbf{r})$  in the Coulomb gauge (i.e.,  $\nabla \cdot \hat{\mathbf{A}} = 0$ ). The photon modes in a given nanophotonic/nanoplasmonic structure can be readily calculated via standard mode decomposition techniques<sup>60</sup>.

In the second quantization, the photonic Hamiltonian can then be written as,

$$\hat{H}_p = \sum_{\alpha} \frac{\hbar\omega}{2} [\hat{a}_{\alpha}\hat{a}_{\alpha}^\dagger + h.c.] = \sum_{\alpha} \hbar\omega (\hat{a}_{\alpha}^\dagger\hat{a}_{\alpha} + 1/2).$$

with the equivalence,

$$A_\alpha \equiv \sqrt{\frac{\hbar}{2\varepsilon_0\omega V}}\hat{a}_{\alpha}, \quad A_\alpha^* \equiv \sqrt{\frac{\hbar}{2\varepsilon_0\omega V}}\hat{a}_{\alpha}^\dagger, \quad (6)$$

Thus, the fields are quantized by the isomorphic association of a quantum mechanical harmonic oscillator to each radiation mode  $\alpha$ . Introducing the canonical position and momentum operators as,  $\hat{q}_{\alpha} = \sqrt{\frac{\hbar}{2\omega_{\alpha}}}(\hat{a}_{\alpha}^\dagger + \hat{a}_{\alpha})$ ,  $\hat{p}_{\alpha} = i\sqrt{\frac{\hbar\omega_{\alpha}}{2}}(\hat{a}_{\alpha}^\dagger - \hat{a}_{\alpha})$ . The photonic Hamiltonian can now be rewritten as,

$$\hat{H}_p = \frac{1}{2} \sum_{\alpha} (\hat{p}_{\alpha}^2 + \omega^2 \hat{q}_{\alpha}^2). \quad (7)$$

Then the full light-matter Hamiltonian in Coulomb gauge can now be fully constructed and written as,

$$\hat{H}_c = \sum_i^{N_e+N_n} \frac{1}{2m_i} (\hat{p}_i - z_i \hat{\mathbf{A}}(\mathbf{r}_i))^2 + \hat{V} + \hat{H}_p, \quad (8)$$

which is often referred to as the minimal coupling Hamiltonian.

We now aim to apply a unitary transformation on Eq. 8 to achieve an expression where the momenta of the molecular DOFs ( $\hat{p}_i$ ) are decoupled from the vector potential ( $\hat{\mathbf{A}}(\mathbf{r}_i)$ ). In other words, we will shift the light-matter coupling from momentum fluctuations into displacement fluctuations. This transformation, referred to as the PZW transformation, can be written as  $\hat{U}_{\text{PZW}} = \exp[-\frac{i}{\hbar}\hat{\mathbf{D}} \cdot \hat{\mathbf{A}}]$  where  $\hat{\mathbf{D}} = \sum_i^{N_n} z_i \hat{\mathbf{R}}_i - \sum_i^{N_e} e \hat{\mathbf{r}}_i$  is the molecular dipole moment. The PZW transformation  $\hat{U}(\mathbf{d})\hat{H}_c\hat{U}^\dagger(\mathbf{d})$  is nothing but a reduction in matter momentum such that  $\hat{p}_i - z_i \mathbf{A} \rightarrow \hat{p}_i$  and a boost in photonic momentum by  $\hat{p}_{\alpha} \rightarrow \hat{p}_{\alpha} + \sqrt{2\omega/\hbar d} \cdot \mathbf{A}_0$ . Applying PZW transformation results in the QED Hamiltonian, referred to as the dipole gauge Hamiltonian,

$$\hat{H} = \hat{T} + \hat{V} + \hat{H}_{ep} \quad (9)$$

where

$$\begin{aligned} \hat{H}_{ep} &= \hat{U}_{\text{PZW}}^\dagger \hat{H}_p \hat{U}_{\text{PZW}} = \frac{1}{2} \sum_{\alpha} [(\hat{p}_{\alpha} + \boldsymbol{\lambda}_{\alpha} \cdot \mathbf{D})^2 + \omega_{\alpha}^2 (\hat{q}_{\alpha})^2] \quad (10) \\ &= \frac{1}{2} \sum_{\alpha} \left[ \hat{p}_{\alpha}^2 + \omega_{\alpha}^2 (\hat{q}_{\alpha} - \frac{\boldsymbol{\lambda}_{\alpha}}{\omega_{\alpha}} \cdot \mathbf{D})^2 \right]. \end{aligned}$$

The second line is reached via canonical transformation between coordinate and momentum operators,  $\hat{p} \rightarrow -\omega \hat{q}$ ,  $\hat{q} \rightarrow 1/\omega \hat{p}$ . Here,  $\boldsymbol{\lambda}_{\alpha} = \sqrt{\frac{1}{\varepsilon V}} \mathbf{u}_{\alpha}$  and  $g_{\alpha} \equiv \boldsymbol{\lambda}_{\alpha} \cdot \mathbf{D}$  defines the light-matter coupling strength, which depends on the volume of quantized photon  $V$  in

radiation mode  $\alpha$ . The total light-matter Hamiltonian can now be *re-partitioned* after the unitary transformation as,

$$\hat{H}_{\text{PF}} = \hat{H}_M + \hat{H}_p + \hat{H}_{ep} + \hat{H}_{\text{DSE}} \quad (11)$$

$$= \hat{H}_M + \sum_{\alpha} \left[ \omega_{\alpha} (\hat{a}_{\alpha}^\dagger \hat{a}_{\alpha} + \frac{1}{2}) + \sqrt{\frac{\omega_{\alpha}}{2}} \boldsymbol{\lambda}_{\alpha} \cdot \hat{\mathbf{D}} (\hat{a}_{\alpha}^\dagger + \hat{a}_{\alpha}) + \frac{1}{2} (\boldsymbol{\lambda}_{\alpha} \cdot \hat{\mathbf{D}})^2 \right]. \quad (12)$$

Here,  $\hat{H}_M = \hat{T}_n + \hat{T}_e + \hat{V}$  is the bare molecular Hamiltonian which includes all Coulomb interactions  $\hat{V}$  between electrons and nuclei as well as the kinetic energy operators of both,  $\hat{T}_e$  and  $\hat{T}_n$ , respectively. This Hamiltonian is often referred to as the Pauli-Fierz (PF) Hamiltonian.

Now it is clear that the light-matter coupling strength is determined by the two quantities: the molecular dipole strength and the quantized cavity volume. Hence, there are two general strategies to enter the strong coupling regime: (I) reduce the cavity volume and (II) increase the number of molecules (increasing the total dipole moment). Therefore, there are two major experimental nanocavity designs for strong light-matter coupling. The first is the nanophotonic cavity that leverages a large number of molecules to achieve strong coupling. The other is the nanoplasmonic cavity that leverages a locally enhanced electric field confined in a small volume to enhance the coupling. This local field is effectively confined to the nearby surrounding of a spherical nanoparticle residing on a lattice of nanoparticles on the scale of nm<sup>3</sup> or even Å<sup>3</sup> (namely picocavities<sup>23,61,62</sup>).

The derivation of Eq. 10 assumes dipole approximation. However, the dipole approximation may fail in the ultra-confined nanoplasmonic cavities where the EM fields are confined within nanometric volumes<sup>63,64</sup>. Consequently, the size of molecules becomes comparable to the cavity volumes, and the widely used dipole approximation breaks down. Position-dependent coupling strength that requires the spatial distribution of excitonic and photonic quantum states is found to be a key aspect in determining the dynamics in ultrasmall cavities both in the weak and strong coupling regimes<sup>63</sup>.

## 2.2 Weak and strong coupling

The light-matter interactions in nanoplasmonic environments can be split into weak and strong coupling regimes. The weak-coupling regime is associated with the Purcell enhancement of spontaneous emission. This effect has been found to be particularly strong when the molecule is placed next to a metallic surface or nanostructure<sup>65</sup>. In such a regime, the plasmon has been found to be able to tune photophysics and photochemistry (i.e., plasmon chemistry) via various possible pathways.

The other regime is the strong coupling regime, where the light-matter interaction cannot be treated perturbatively. The strong coupling is characterized by a reversible (coherent) exchange of energy (known as Rabi oscillation) between the matter and the cavity photon, as the coupling is strong enough to compete with the dissipation. In this regime, the formation of polariton requires theoretical methods that treat the matter and photonic DOFs on equal footing and describe the multiple interac-

tions between photons, electrons, and nuclei on different length and time scales (polariton chemistry).

### 3 Plasmon chemistry

Although confined plasmonic modes result in localized and enhanced fields, plasmons usually suffer from strong dissipation, making it hard to enter the strong-coupling regime in plasmonic systems. Nevertheless, even without strong coupling, plasmonics provides a unique setting for manipulating light via the confinement of EM (below the diffraction limit). Such extreme concentration of EM field<sup>66</sup> has led to a wide range of applications, such as plasmon-enhanced molecular spectroscopy<sup>8,67–70</sup>, photovoltaics<sup>71,72</sup>, nanophotonic lasers and amplifiers<sup>73–76</sup>, quantum information<sup>16</sup>, and many others<sup>18,77,78</sup>.

Following the ultrafast plasmon excitation, nonradiative plasmon decay leads to the formation of energetic electron–hole pairs (namely hot-carriers)<sup>79–83</sup>, which are highly nonthermal and can have considerably higher energies than those rising from thermal equilibrium. The hot electrons (HEs) (and their corresponding holes) redistribute their energies quickly as a result of electron–electron scattering<sup>79,80</sup>, reaching a quasi-thermal equilibrium but with a high effective temperature. Further cooling of the hot electrons takes place via energy dissipation into the phonon modes of the nanoparticle, and the energy is ultimately dissipated to the surroundings via thermal conduction. Nevertheless, investigations in the last two decades have found that chemical reactions can be stimulated by localized EM fields (or photons), and electronic and/or thermal energies (that result from plasmon decay) via various different pathways (more details in Sec. 3.3). This leads to an emerging field of plasmon chemistry that designs nanostructure-based surface plasmons as mediators to redistribute and convert photon energy in various time, space, and energy scales to drive chemical reactions<sup>20–22,70,84–95</sup>.

In most scenarios, the nanoplasmionic properties are obtained by solving Maxwell's equations with proper dielectric function and boundary conditions. However, Maxwell's equations fail to describe the quantum effects (such as the tunneling in charge transfer mode and quantum confinement<sup>96</sup>), and quantum theory for plasmon (quantum plasmonics) is required in nanoscale.

#### 3.1 Ab initio method for quantum plasmonics

From a quantum mechanical point of view, a plasmon is nothing but a specific collective excitation. Hence, the quantum chemistry method for excited states can be used to predict the optical properties of plasmonic nanostructures<sup>97</sup>. The many-body perturbation theory, time-dependent density functional theory (TDDFT), or time-dependent density functional tight-binding (TDDFTB) are methods of choice to compute the plasmon excitations<sup>98–106</sup>. Without losing generality, here we use TDDFT theory as an example to demonstrate the computation of plasmon excitations.

TDDFT is the formal extension of the Hohenberg-Kohn-Sham density functional theory (DFT). Within the DFT, The Hohenberg-Kohn (HK) theorem<sup>107</sup> states that the ground-state electron density unambiguously defines the many-electron ground state for an  $N$ –electron system under the influence of an external potential.

And there exists an energy functional that guarantees the ground state energy can be reached by the variational principle though the exact functional is an unsolved problem,

$$E[\rho] = T + V_H(\mathbf{r}) + E_{xc}[\rho] + \int V(\mathbf{r})\rho(\mathbf{r})d\mathbf{r}. \quad (13)$$

Where  $V_H[\rho] = \int \frac{\rho(\mathbf{r}')}{|\mathbf{r}-\mathbf{r}'|} d\mathbf{r}'$  is the Hartree potential,  $E_{xc}$  is the exchange-correlation functional, which contributes < 10% to the total energy, but describes the most critical correlation effects<sup>108</sup>. But DFT is a ground-state theory. Light-matter interactions usually result in many elementary excitations within the matter. Among various many-body methods, TDDFT (the formal extension of DFT theory) is the preferred tool to evaluate the excited states and optical properties in extended molecular or condensed matter systems.

In the linear response regime, the induced electron density due to the light-matter interaction  $\delta V(\mathbf{r})$  is given by

$$\delta\rho(\mathbf{r}) = \int \chi(\mathbf{r}, \mathbf{r}') \delta V_{ext}(\mathbf{r}') d\mathbf{r}'. \quad (14)$$

Where  $\chi(\mathbf{r}, \mathbf{r}')$  is the polarizability describing the density response of the many-body ground state with respect to an external perturbation. In addition, according to Runge-Gross theorem<sup>109</sup>,  $\delta\rho(\mathbf{r})$  is also the induced density of the KS system, but due to a perturbation

$$\delta\rho(\mathbf{r}) = \int \chi_{KS}(\mathbf{r}, \mathbf{r}') \delta V_{KS}(\mathbf{r}') d\mathbf{r}'. \quad (15)$$

$\chi_{KS}$  here is the linear response of the KS electrons, which can be trivially evaluated from the KS orbitals and energies.

$$\delta V_{KS}(\mathbf{r}) = \delta V_{ext}(\mathbf{r}) + V_H[\delta\rho(\mathbf{r})] + \delta V_{xc}(\mathbf{r}). \quad (16)$$

The induced XC potential is given by  $\delta V_{xc}(\mathbf{r}) = \int f_{xc}(\mathbf{r}, \mathbf{r}') \delta\rho(\mathbf{r}') d\mathbf{r}'$ , where  $f_{xc}$  is the dynamical XC kernel  $\frac{\delta V_{xc}(\mathbf{r})}{\delta n(\mathbf{r}')}$ <sup>110</sup>,

$$f_{xc}(\mathbf{r}, \mathbf{r}') = \frac{\delta V_{xc}(\mathbf{r})}{\delta\rho(\mathbf{r}')}, \quad (17)$$

Hence, a linear equation for the induced density can be derived from Equations 14-17,

$$[1 - \chi_{KS}(\omega) f_{hxc}(\omega)] \chi(\omega) \delta V_{ext}(\omega) = \chi_{KS}(\omega) \delta V_{ext}(\omega). \quad (18)$$

where  $f_{hxc}(\mathbf{r}, \mathbf{r}') = \frac{1}{|\mathbf{r}-\mathbf{r}'|} + f_{XC}(\mathbf{r}, \mathbf{r}')$ . Casting Equation 18 into the matrix of coupled KS single excitations, it is possible to calculate the excitation energies  $\omega_s$  of the system (the poles of the response function) and transition densities (and oscillator strengths). Consequently, the response function  $\chi(\omega)$  can be rewritten as

$$\chi(\mathbf{r}, \mathbf{r}', \omega) = 2 \sum_s \rho_s(\mathbf{r}) \rho_s(\mathbf{r}') \zeta_s(\omega), \quad (19)$$

where  $\zeta_s(\omega) = \frac{1}{\omega - \omega_s + i\eta} - \frac{1}{\omega + \omega_s - i\eta}$  and  $\eta$  represents a positive infinitesimal. The factor of 2 comes from summation over spin indices. The excitation energies in the system are denoted as  $\omega_s$ , which are calculated from the Casida method<sup>110</sup>.  $\rho_s(\mathbf{r})$  is the transition density, which can be expanded in terms of electronic

transitions between occupied state  $i$  to unoccupied states  $a^{111}$ ,

$$\rho_s(\mathbf{r}) = \sum_{ia} X_{ia}^s \psi_i(\mathbf{r}) \psi_a(\mathbf{r}) \left( \frac{\epsilon_a - \epsilon_i}{\omega_s} \right)^{1/2}, \quad (20)$$

where  $\epsilon_i$  are the KS eigenvalues and the corresponding molecular orbitals are  $\psi_i$ . And  $X_{ia}^s$  are the Casida transition coefficient from the occupied  $i$  state to the unoccupied  $a$  state of the  $s^{th}$  excitation. By examining the nature of  $X_{ia}^s$ , it is possible to distinguish between normal excitation and plasmonic excitation. Plasmonic excitations generally are characterized by collective transitions from occupied to virtual orbitals<sup>112,113</sup>.

### 3.2 Hot electron generation and relaxation

Once the plasmon excitation is obtained, the electron-plasmon interaction can be described by the Hamiltonian

$$\hat{H}_{int} = \frac{e}{2m_e} \int d\mathbf{r} \hat{\Psi}^\dagger V_{eff}(\mathbf{r}) \hat{\Psi}, \quad (21)$$

where  $V_{eff}(\mathbf{r})$  is the effective potential induced by the excitation, which can be calculated from the transition density  $\rho_s(\mathbf{r})$  by following Lundqvist's approach<sup>111,112</sup>. The polarizability of the NP upon external excitation is given by Equation 19. And the induced potential is given by<sup>111</sup>,

$$V_{eff}(\mathbf{r}) = \epsilon^{-1}(\mathbf{r}, \mathbf{r}') \delta V_{ext}(\mathbf{r}'). \quad (22)$$

Where the dielectric function  $\epsilon^{-1}(\mathbf{r}, \mathbf{r}')$  is given by

$$\epsilon^{-1}(\mathbf{r}, \mathbf{r}') = \delta(\mathbf{r}, \mathbf{r}') + \int d\mathbf{r}_1 f_{hxc}(\mathbf{r}, \mathbf{r}_1) \chi(\mathbf{r}_1, \mathbf{r}', \omega). \quad (23)$$

Thus, within the second quantization, electron-excitation Hamiltonian  $\hat{H}_{eps}$  reads

$$\hat{H}_{eps} = \sum_{ij} \left[ \mathcal{M}_{ij}^s \hat{c}_i^\dagger \hat{c}_j \hat{b}_s + \text{h.c.} \right], \quad (24)$$

where  $\mathcal{M}_{ij}^s$  is the electron-excitation coupling strength, describing the scattering of quasiparticles from state  $i$  into state  $j$  via the emission or absorption of excitation in the state  $s$ . These elements are given by (see SI for details)

$$\mathcal{M}_{ij}^s = \zeta_s(\omega) \mathbf{E} \cdot \mathbf{f}^s \langle \psi_i(\mathbf{r}_1) | V_H^s(\mathbf{r}_1) | \psi_j(\mathbf{r}_1) \rangle \quad (25)$$

where  $\mathbf{f}^s = \sqrt{\frac{2m_e \omega_s}{2\hbar^2}} \sum_{ia} X_{ia}^s \left( \frac{\epsilon_a - \epsilon_i}{\omega_s} \right)^{1/2} \mathbf{d}_{ia}$  is the oscillator vector and  $|\mathbf{f}^s|^2 = \frac{2m_e}{3\hbar^2} \omega_s |\langle \Psi_0 | \mathbf{r} | \Psi_s \rangle|^2$  is the oscillator strength.  $V_H^s(\mathbf{r}) = \int d\mathbf{r}' \frac{\rho_s(\mathbf{r}')}{|\mathbf{r} - \mathbf{r}'|}$  is the Hartree potential induced by the transition density. Detailed derivation can be found in the SI. Above equation actually includes both plasmon excitation ( $\zeta_s(\omega) \mathbf{E} \cdot \mathbf{f}^s$ ), screening effect and hot electron-hole pairs generation ( $\langle \psi_i(\mathbf{r}_1) | V_H^s(\mathbf{r}_1) | \psi_j(\mathbf{r}_1) \rangle$ ). Neglecting the screening effect would result in a much larger electron-plasmon coupling matrix and shorter HC lifetime distribution<sup>114</sup>. In contrast, our model connects the widely used semiempirical model and the recently developed quantum model with the plasmon excitation and HC generation treated on equal footing. Equation 24 is the general formalism that describes the coupling between electrons and photo-

toexcitation. If the external field matches the plasmon energy, plasmon resonance will be excited, and Equation 24 reduces to the electron-plasmon coupling.

**HC generation.** The electron-plasmon coupling describes the HC generation following plasmon decay. After the electron-plasmon coupling matrix is obtained, the HC generation can be readily calculated from the Fermi golden rule<sup>79,83</sup>,

$$\Gamma_{i \rightarrow a}^{ex} = \frac{4}{\hbar} \sum_s |\mathcal{M}_{ia}^s|^2 \frac{\gamma_{ex}}{(\epsilon_i - \epsilon_a + \omega_s)^2 + \gamma_{ex}^2} \delta(\omega - \omega_s). \quad (26)$$

$\gamma_{ex}$  refers to the linewidth of the excitation.  $\omega$  is the excitation energy.  $\delta(\omega - \omega_s)$  describes the generalized photoexcitation. Hence, Equation 26 describes the photoexcitation and HC generation. When the energy of the external field  $\omega$  matches the plasmon energy, Equation 26 describes the HC generation from plasmon decay. Otherwise, it describes the HC generation from regular excitation.

**HC relaxation.** After generation from plasmon decay, the HCs will undergo a relaxation process mainly due to the electron-electron and electron-phonon scatterings. The dynamics of the HCs are investigated by propagating the density matrix  $\rho(t)$ . The equation of motion (EOM) of  $\rho$  is subject to the following quantum Liouville Von-Neumann equation<sup>115</sup>,

$$i\partial_t \rho(t) = i\partial_t \rho(t)|_{coh} + \mathcal{L}_{ee}[t] + \mathcal{L}_{ph}[t] + \mathcal{L}_s[t] \quad (27)$$

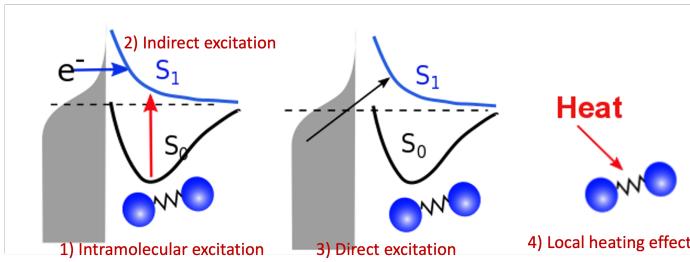
The first part on the right-hand side (RHS) of the above equation describes the coherent evolution of the density matrix between different states. The second and third parts on the RHS of Equation 27 represent the dissipations induced by the electron-electron and electron-phonon scattering effects, respectively. Finally, the last term on the RHS of Equation 27 described the extraction of HCs. In general, in the presence of electron-electron and electron-phonon interaction, the dissipation can be described by a Liouville equation by employing the many-body perturbation theory (MBPT)<sup>116</sup>,  $\mathcal{Q}(t) = \mathcal{D}(t) - \mathcal{D}^\dagger(t)$ . Where the dissipation matrix  $\mathcal{D}_t$  can be written in terms of Green's functions  $G$  and self-energies  $\Sigma$ <sup>79,116</sup>, i.e.,

$$\mathcal{D}(t) = i \int^t dt'_{-\infty} [\Sigma^{<}(t, t') G^{>}(t', t) - \Sigma^{>}(t, t) G^{<}(t', t)]. \quad (28)$$

Hence, the key point is to develop approximations to the self-energies  $\Sigma^{<,>}(t, t')$  and efficient numerical methods to compute the  $\mathcal{D}(t)$ . The detailed formalisms of  $\mathcal{D}(t)$  for electron-electron and electron-phonon scatterings can be found in Ref.<sup>79</sup>. But it should be noted that the approximation used for electron-electron scattering should conserve both particles and energy.

### 3.3 Plasmon-mediated chemical reactivities and theoretical challenges

However, the computations of plasmon excitation and distribution of hot electrons (from plasmon decay) are insufficient to give insights into plasmon-mediated chemistry because plasmon-induced photocatalysis is a complex dynamical process that involves multiple interactions and mechanisms. To date, four major microscopic mechanisms have been proposed to explain how



**Fig. 2** (a) Proposed mechanisms of plasmon-mediated reactivities. 1) Enhanced intramolecular excitation, 2) direct charge transfer, 3) indirect charge transfer, and 4) local heating.

plasmonics facilitates various chemical reactions through the concentration of light and hot-carrier dynamics (Figure. 2). The first mechanism involves the extreme concentration of light, which significantly promotes the excitation of electrons within the adsorbed molecule. Such an excitation takes place within the adsorbate only but is significantly enhanced by plasmonics, which is referred to as the enhanced intramolecular excitation mechanism<sup>90,92,117</sup>. The second mechanisms involve the transfer of plasmonic hot carriers. Due to the hybridization of the adsorbate–nanoparticle system, the generated hot electrons (or holes) can transfer to the adsorbed molecules. Such transfers result in the placement of an adsorbate–nanoparticle system onto a manifold of excited potential energy surfaces (PESs), where the adsorbed molecule experiences strong forces that activate its internal vibrational excitations and, ultimately, chemical transformation. This mechanism is attributed to an indirect HE transfer from metal nanoparticles to the adsorbed molecule<sup>22,91</sup>. Third, due to the hybridization, direct excitation of electrons from metal states near the Fermi level to the unoccupied molecular orbital (LUMO) of the adsorbed molecule can occur when the plasmon frequency is resonant with the excitation energy between the metallic occupied orbitals and hydrated metal-molecular orbitals. Such a reaction pathway is referred to as the direct charge transfer (CT) mechanism. Such excitation circumvents the thermalization of HEs<sup>94,118–120</sup>, but requires matching between the plasmon energy and the energy gap between the metal states and the molecular LUMO. Finally, local heating resulting from hot-carrier relaxation can thermally activate a reaction<sup>93,121,122</sup>.

These mechanisms share some common features that render their clear distinction very challenging. Indeed, different mechanisms have been proposed even for the same chemical reaction, leaving a very confusing situation. Hence, in order to obtain a comprehensive understanding of plasmon-mediated chemistry, it's essential for the theoretical methods to fulfill the following requirements:

1. Efficient computation of a dense manifold of excited states because plasmonic excitations in metallic nanostructures are usually not low-lying states. Even for a small nanoparticle (2 nm diameter, for example), thousands of excited states are to be computed to reach the plasmon excitations<sup>87</sup>.
2. The ability to capture the nonadiabatic transition between excited states since the indirect transfer mechanism involves

the transition between the hot electron state (excitations localized within the plasmonic system only) and the charge transfer state<sup>89</sup>.

3. Inclusion of electron-vibrational couplings that lead to the hot electron relaxation and local heating.
4. Inclusion of electron-electron scattering that leads to the redistribution of hot electrons<sup>82,123</sup>.

In particular, recent debates on the thermal impact underpin the necessity of atomistic and dynamical insights via nonadiabatic simulations of the HE generation, transfer, and relaxation processes on equal footing.

### 3.4 Nonadiabatic simulation of Plasmon-mediated chemical reactivities

**Ehrenfest dynamics and Surface Hopping.** With the Born-Oppenheimer approximation, the electronic wave function  $\Theta(\mathbf{r}, \mathbf{R})$  is expanded in the basis of adiabatic BO states, which depend on the electronic coordinates  $\mathbf{r}$  and the nuclear coordinates  $\mathbf{R}(t)$  according to

$$\Theta(\mathbf{r}, \mathbf{R}) = \sum_{n=1}^{N_{st}} c_n(t) |\phi_n(\mathbf{r}, \mathbf{R}(t))\rangle. \quad (29)$$

Here  $N_{st}$  is the total number of adiabatic electronic states,  $|\phi_n(\mathbf{r}, \mathbf{R}(t))\rangle$  is the adiabatic electronic wavefunction of state  $n$  and  $c_n(t)$  are the time-dependent complex expansion coefficients.  $\mathbf{R}(t) = \{\mathbf{R}_A(t)\}_{A=1}^{A=N_A}$  ( $N_A$  is the total number of atoms in the system) are the nuclear trajectories which are obtained by solving the classical Newton's equations of motion (EOMs) with the mixed quantum-classical framework<sup>124</sup>,

$$M_A \frac{d^2 \mathbf{R}_A}{dt^2} = -\nabla_{\mathbf{R}_A} E(\mathbf{R}), \quad (30)$$

where  $M_A$  is the mass of  $A^{th}$  atom.  $E(\mathbf{R})$  is the potential energy surface (PES), which can be an averaged one  $E(\mathbf{R}) = \sum_n C_n(t) E_n(\mathbf{R})$  within the mean-field Ehrenfest dynamics or a single PES of a certain state  $E_k(\mathbf{R})$  within the Surface hopping (SH) framework<sup>124–126</sup>.

By substituting Eq. 29 into the time-dependent Schrödinger equation and keeping only the first-order nonadiabatic coupling terms, a set of EOMs for the coefficients  $c_n(t)$  along a given classical trajectory can be obtained<sup>124,127</sup>

$$i\hbar \frac{\partial c_n(t)}{\partial t} = c_n(t) E_n(\mathbf{R}) - i\hbar \sum_m c_m(t) \dot{\mathbf{R}} \cdot \mathbf{d}_{nm}. \quad (31)$$

Here the orthogonal condition of adiabatic states  $\langle \Psi_n | \Psi_m \rangle = \delta_{nm}$  is used.  $\mathbf{d}_{nm} = \langle \Psi_n | \nabla_{\mathbf{R}} | \Psi_m \rangle$  is the nonadiabatic derivative coupling term (or nonadiabatic coupling vector, NACR). A key variable in Eq. 31 is the time-derivative nonadiabatic coupling scalar (NACT) between two adiabatic states

$$\dot{\mathbf{R}} \cdot \mathbf{d}_{nm} = \langle \Psi_n | \frac{\partial}{\partial t} | \Psi_m \rangle, \quad (32)$$

which is responsible for the nonadiabatic transitions between different adiabatic states and can be easily calculated with many ab

initio methods for excited states.

With the SH algorithm, the state that governs the dynamics of nuclei is determined by a stochastic process. The time-dependent coefficients  $c_n(t)$  obtained in Eq. 31 are used to calculate the hopping probabilities between different electronic excited states within the framework of the FSSH algorithm. The hopping probabilities between excited states  $n$  and  $m$  are given by<sup>128</sup>

$$g_{n \rightarrow m}(\mathbf{R}, t) = \frac{\int_t^{t+N_q \delta t} dt b_{mn}(\mathbf{R}, t)}{a_{mn}(t)}, \quad (33)$$

where  $N_q = \frac{\Delta t}{\delta t}$ , with  $\Delta t$  and  $\delta t$  correspond to the time steps for evolving motions of nuclei and electrons in Eq. 30 and Eq. 31, respectively. The chosen value of  $\delta t$  must be small enough to resolve strongly localized peaks in NACT in order to avoid underestimation of transition probabilities. This is particularly important when crossings between adiabatic states are encountered in the trajectory, in which  $\delta t$  will be further refined.  $a_{nn}(t) = c_n(t)c_n^*(t)$  defines the time-dependent density matrix elements, and  $b_{mn}(\mathbf{R}, t) = -2Re(a_{nm}^* \dot{\mathbf{R}} \cdot \mathbf{d}_{nm})$ . Note that  $g_{n \rightarrow m} = -g_{m \rightarrow n}$  and  $g_{n \rightarrow n} = 0$  since  $\mathbf{d}_{nm}$  are antisymmetric. Hopping between adiabatic states is determined stochastically by comparing  $g_{n \rightarrow m}$  to a random number  $\xi (\xi \in (0, 1))$ . A hop from state  $n$  to state  $m$  is performed if

$$\sum_{l=1}^{m-1} g_{n \rightarrow l} < \xi \leq \sum_{l=1}^m g_{n \rightarrow l}, \quad (34)$$

where states are assumed to be ordered with increasing transition energy. On the other hand, the system remains in state  $n$  when  $\sum_{l=1}^{N_{st}} g_{n \rightarrow l} < \xi < 1$ . If  $g_{n \rightarrow m} < 0$ , the hop is unphysical, and the probability is set to zero. Finally, if a hop to a higher energy state is predicted, there must be sufficient nuclear kinetic energy along the direction of NACR. Otherwise, the hop is rejected. After a successful hop, the total electron-nuclear energy is conserved by rescaling the nuclear velocity in the direction of the NACR according to the procedure described in reference<sup>126,129</sup>. In addition, during the dynamics, we monitor the relative phase of the ground to excited state transitions and maintain the same phase (sign) to avoid a sudden sign change in the NACT. This is done by enforcing the sign of the largest component of the Casida eigenvectors to the same along the trajectory.

Despite broad popularity in the community, either Ehrenfest dynamics or the surface hopping (SH) approach have well-known limitations such as generating artificial electronic coherence or giving incorrect long-time population<sup>126</sup>. To address these challenges and provide accurate dynamics, a multiconfigurational Ehrenfest (MCE) dynamics approach<sup>130</sup> and ab initio multiple cloning (AIMC)<sup>131–134</sup> are developed accordingly.

**Multiconfigurational Ehrenfest (MCE).** MCE generalizes EHR formalism by representing the wave function as a linear combination of Ehrenfest configurations. Each MCE configuration moves along its own Ehrenfest (mean-field) trajectory. Within the MCE formalism, the molecular wavefunction  $|\Psi\rangle$  is expressed in the trajectory-guided Gaussian basis functions (TBF) representation ( $|\psi_n\rangle$ ),

$$|\Psi(t)\rangle = \sum c_n |\psi_n(t)\rangle. \quad (35)$$

And each configuration (or TBF) is described by the product of nuclear and electronic parts,

$$|\psi_n(t)\rangle = |\chi_n(\mathbf{R}, t)\rangle \sum_I a_I^n |\phi_I^n(\mathbf{r}, \mathbf{R}(t))\rangle. \quad (36)$$

Where  $|\phi_I^n(\mathbf{r}, \mathbf{R}(t))\rangle$  is the adiabatic state of configuraiton  $n$ .  $|\chi_n\rangle$  are Gaussian nuclear basis functions

$$|\chi_n\rangle = \left( \frac{2\alpha}{\pi} \right)^{N_d/4} e^{\{-\alpha(R-\bar{R}) + \frac{i}{\hbar}P(R-\bar{R}) + \frac{i}{\hbar}\gamma_n(t)\}}. \quad (37)$$

The couplings between TBFs in the MCE approach are described by the EOM of  $c_n(t)$ , which can be readily obtained by substituting Eq. 35 into the Schrödinger equation:

$$i\hbar \sum_n S_{mn} \dot{c}_n = \sum_n \left[ H_{mn} - i\hbar \langle \psi_m | \frac{d\psi_m}{dt} \rangle \right] c_n. \quad (38)$$

where

$$H_{mn} = \sum_{I,J} (a_I^m)^* a_J^n \langle \chi_m \phi_I^m | T + V | \chi_n \phi_J^n \rangle, \quad (39)$$

and the overlap  $S_{mn}$  is

$$S_{mn} = \langle \psi_m | \psi_n \rangle = \langle \chi_m | \chi_n \rangle \sum_{I,J} (a_I^m)^* a_J^n \langle \phi_I^m | \phi_J^n \rangle. \quad (40)$$

The nuclear part of Eq. 39 can be obtained analytically,

$$\langle \chi_m \phi_I^m | V | \chi_n \phi_J^n \rangle = \langle \chi_m | -\frac{\hbar^2}{2} \nabla_{\mathbf{R}} M^{-1} \nabla_{\mathbf{R}} | \chi_n \rangle \langle \phi_I^m | \phi_J^n \rangle. \quad (41)$$

While the electronic part (or the potential energy matrix elements) are approximated by<sup>134</sup>

$$\begin{aligned} \langle \chi_m \phi_I^m | V | \chi_n \phi_J^n \rangle &= \frac{1}{2} \langle \phi_I^m | \phi_J^n \rangle \langle \chi_m | \chi_n \rangle \times \\ &\left\{ (V_I^m + V_J^n) + \frac{i}{4\alpha\hbar} (\mathbf{P}_n - \hat{P}_m) \cdot (\nabla_{\mathbf{R}} V_I^m + \nabla_{\mathbf{R}} V_J^n) \right. \\ &\left. - \frac{1}{2} (\mathbf{R}_m - \mathbf{R}_n) \cdot (\nabla_{\mathbf{R}} V_I^m - \nabla_{\mathbf{R}} V_J^n) \right\}. \end{aligned} \quad (42)$$

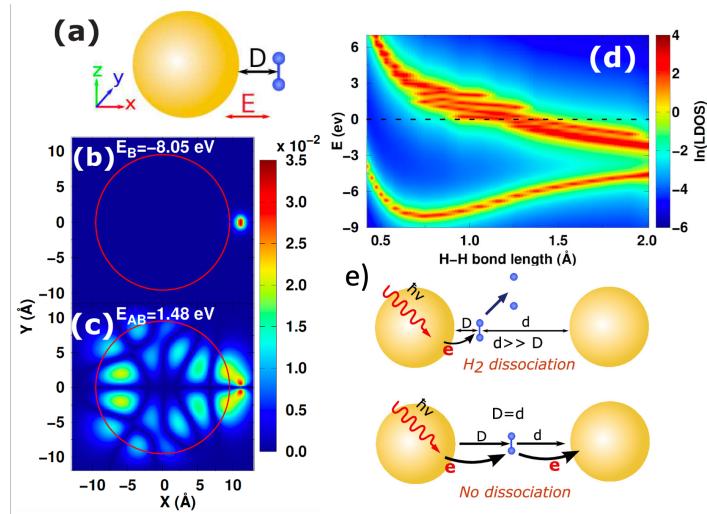
As shown above, within the MCE schemes, the electronic states are now different for different configurations. The overlaps between TBFs have to be calculated and taken into account.

**Ab initio multiple cloning (AIMC).** The AIMC method combines the best features of ab initio Multiple Spawning (AIMS)<sup>135</sup> and Multiconfigurational Ehrenfest (MCE) methods. Similar to the MCE method, the individual trajectory basis functions (TBFs) of AIMC follow Ehrenfest equations of motion. However, the basis set is expanded in a similar manner to AIMS when these TBFs become sufficiently mixed. Consequently, AIMC avoids prolonged evolution on the mean-field potential energy surface (PES).

Within the MCE formalism, the Ehrenfest basis set is guided by an average potential, which is accurate for dynamical processes where the coupling between states persists in time between nearly parallel PES. But the mean-field treatment can be unphysical when the PES of two or more populated electronic states become different in shape, which leads to wave packet branching after leaving the nonadiabatic coupling region. To deal with these cases, the AIMC algorithm is applied to expand the original

basis set of TBFs by "cloning" one TBF into two copies in a way that does not alter the original wave function<sup>131–133</sup>. This is done by creating one of the clones  $|\psi_{n_1}\rangle$  in a pure state and the other clone  $|\psi_{n_2}\rangle$ , which includes contributions from all other electronic states. The corresponding MCE amplitudes  $\{c_{n_1}, c_{n_2}\}$  are adjusted to conserve the original wavefunction<sup>133,134</sup>.

It should be noted the MQC method can be complemented with any electronic structure solvers as long as the gradients and NACs are available. Practical applications need to balance accuracy and numerical efficiency.



**Fig. 3** (a) Schematic diagram of  $\text{H}_2$  molecule adsorbed on the NP surface. b)-c) Spatial distribution of the bonding and antibonding orbitals of  $\text{H}_2$  on Jellium NP. d) Local DOS of bonding and antibonding orbitals of  $\text{H}_2$  modules as a function of H-H bond length. e) Depending on the symmetry, the  $\text{H}_2$  dissociation can be suppressed or restored in the plasmonic dimer. Figures are adapted with permission from Ref. 88. Copyright 2018 American Chemical Society.

### 3.5 Simulation of plasmon-mediated phenomena

#### 3.5.1 Jellium model-based NAMD simulation

Compared to the costly atomistic ab initio calculations for plasmonic nanoparticles, the optical absorption of simple sp-metal nanoparticles described by the Jellium model can be easily calculated and analyzed<sup>83,88,97,136–138</sup>. In general, good agreement with experimental optical properties can be achieved by tuning the Jellium radius. Using the Jellium model, we investigated the generation and relaxation of plasmonic hot carriers<sup>79</sup>, as well as the atomic-scale mechanism of plasmonic hot-carrier-mediated chemical processes such as  $\text{H}_2$  dissociation<sup>88</sup>. Our numerical simulations showed that after photoexcitation, hot carriers transfer to the antibonding state of the  $\text{H}_2$  molecule from the nanoparticle, leading to a repulsive-potential-energy surface and  $\text{H}_2$  dissociation (Figure ??(b-d)). This process occurs when the molecule is close to a single nanoparticle. However, in a plasmonic dimer, dissociation can be inhibited due to sequential charge transfer that effectively reduces the occupation of the antibonding state, as shown in Figure 3(e). When the molecule is asymmetrically positioned in the gap, the symmetry is broken, and dissociation

is restored by significantly suppressing additional charge transfer. Thus, these models illustrate the potential for structurally adjustable photochemistry through plasmonic hot carriers.

#### 3.5.2 TDDFT calculations.

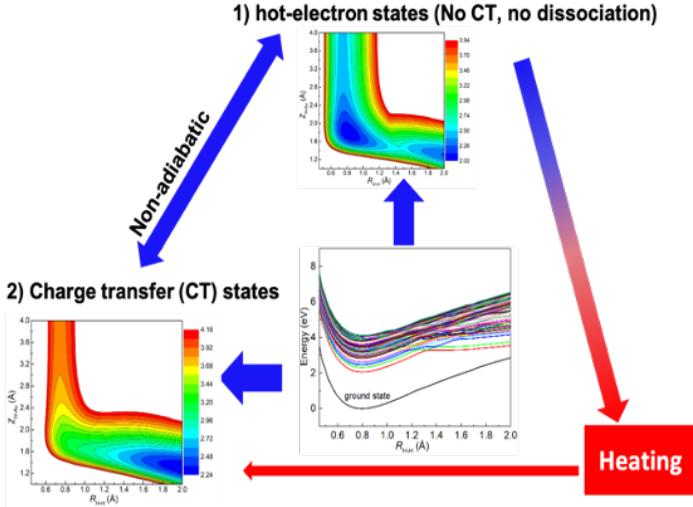
However, the Jellium model oversimplifies the electronic structures of the plasmonic nanostructures, and it lacks atomistic details. Insights from PESs built on the ab initio atomistic model are essential for a more in-depth understanding of plasmon chemistry. Thus, we employed the linear response TDDFT (LR-TDDFT) calculations within the Casida formalism to compute the adiabatic PESs of the  $\text{H}_2$  molecule adsorbed on an Au6 cluster ( $\text{H}_2@\text{Au6}$ )<sup>89</sup>, in order to explore key pathways in LSPR-promoted chemical reactions. Despite the model system is too small to support plasmonic mode and thus cannot describe the dephasing of plasmons that produce hot electrons, the key point of using the model system is to capture key aspects of the later stages of plasmon-facilitated photocatalysis, thus providing mechanistic insights.

Our findings based on DFT calculations indicate that in the ground state, the Au6 cluster supports the adsorption of the  $\text{H}_2$  molecule at the tip site. We further use LR-TDDFT calculations within the Casida formalism to determine the adiabatic excited states and corresponding oscillator strengths. We calculate two-dimensional PESs in the desorption and dissociation reaction coordinates and observe that the adiabatic excited PESs bear similarity to the ground state PES in the Franck-Condon region. Therefore, the  $\text{H}_2$  adsorbate is comparatively stable, as corroborated by the relatively low dissociation and desorption probabilities from our quantum dynamics simulations. By developing the orbital wave function overlapping (OWO) diabatization scheme, we are able to divide the dense manifold of the excited state into two groups: 1) one group is dominated by electronic excitations confined to the Au6 cluster, which can be likened to HE states in metal nanoclusters; 2) the other group of excited states has the antibonding  $\sigma^*$  characteristics of the  $\text{H}_2$  adsorbate due to the hybridization between  $\text{H}_2$  and Au6 cluster, which are denoted as CT states. The crossings among the HE and CT states provide pathways leading from the excited HE states to CT states via nonadiabatic transitions. Quantum dynamics simulations on the diabatic PESs demonstrate that the CT diabatic states are able to drive  $\text{H}_2$  dissociation efficiently and thus are responsible for the experimentally observed HD formation on Au nanoparticles. Our results nevertheless give a clear physical picture of photoinduced  $\text{H}_2$  dissociation on Au clusters.

The presence of HE-CT crossings is not unique to plasmonic catalysis. Such features have also been found in other nonplasmonic catalysis, such as those on the surface of semiconductors<sup>139</sup>. However, plasmonic materials are unique in generating a high concentration of HE due to their larger absorption cross sections. The combination of a high concentration of HE and HE-CT crossings is thus the distinctive feature of plasmonic catalysis.

#### 3.5.3 TDDFTB-based NAMD simulations

As shown in Figure 4, plasmon-mediated chemistry involves a dense manifold of excited states, nonadiabatic transitions be-



**Fig. 4** Schematic diagram of competition among different pathways in plasmon-mediated chemical reactions on a dense manifold of excited states. The excited states can be divided into HE and CT states. The CT states are responsible for the chemical reaction, which can be triggered by the nonadiabatic transition between HE and CT states. Such nonadiabatic transitions have to compete with hot carrier relaxations that lead to local heating.

tween HE and CT states, and their competition with various dissipation channels. Such complicated processes require nonadiabatic simulations that treat the HE generation, relaxation, and HE-CT transitions treated on equal footing. To this end, larger clusters should be used in modeling plasmonic catalysis, and more efficient semiempirical methods, such as those based on time-dependent density functional tight binding (TDDFTB)<sup>140</sup>, might be needed for the efficient electronic structure calculations. To this aim, We recently developed an efficient NAMD method by combing the TSH algorithm and LR-TDDFTB method.<sup>128,141–143</sup> LR-TDDFTB is the tight-binding extension of TDDFT,<sup>141,144</sup> which has been successfully applied to investigate the optical properties of plasmonic NPs.<sup>145,146</sup> It enables calculations of plasmonic excitation along with several hundred excitation states per time step. The relaxation processes of plasmon-induced by electron-phonon interactions are treated by the TSH algorithm. With the NAMD-DFTB method, we demonstrate the plasmon relaxation of an Au<sub>20</sub> cluster. Our simulations show that Au<sub>20</sub> can support plasmon-like excitation, which includes the superposition of multiple single-particle excitation components.

The numerically efficient LR-TDDFTB method allows us to address a dense manifold of excited states to ensure the inclusion of plasmon excitation. Starting from the photoexcited plasmon states in Au<sub>20</sub> cluster, we find that the time constant for relaxation from plasmon excited states to the lowest excited states is about 2.7 ps, mainly resulting from a step-wise decay process caused by low-frequency phonons of the Au<sub>20</sub> cluster. Furthermore, our simulations show that the lifetime of the phonon-induced plasmon dephasing is ~10.4 fs, and such a swift process can be attributed to the strong nonadiabatic effect in small clusters. Our simulations demonstrate a detailed description of the dynamic processes in nanoclusters, including plasmon excitation, hot carrier generation from the plasmon excitation dephasing, and the subsequent

phonon-induced relaxation process.

The NEXMD-DFTB method was employed to investigate the plasmon-induced bond activation of CO adsorbed on Au<sub>20</sub> cluster<sup>87</sup>. The simulations provide a comprehensive and accurate depiction of the multiple dynamic processes involved in plasmon-mediated chemistry, including plasmon excitation, HEs relaxation, direct/indirect HE transfer, and the activation of CO vibration mode induced by HE transfers. These simulations reveal the critical role of charge transfer (CT) states in plasmon-induced CO activation. The CT states excite both direct and indirect HE transfer, which leads to the activation of CO stretching mode, an essential component of plasmon energy relaxation. Our simulations demonstrate the high efficiency of CO vibrational mode activation, achieving a success rate of approximately 40%. Notably, the HE transfer occurs at a faster rate than the conventional scattering process of the Au<sub>20</sub>, completing within approximately 100 fs, while the energy relaxation takes place over a timescale of approximately 1 ps. Furthermore, our direct atomistic simulations provide detailed insights into the potential energy evolution during the plasmon-mediated chemical transformation, thereby enabling a comprehensive understanding of the energy relaxation and HE transfers during the reaction, as well as elucidating the reaction pathways.

## 4 Polariton chemistry

### 4.1 Introduction and background

When the coupling of the light and molecular DOFs is strong compared to the dissipation of the excited state population (*e.g.*, cavity leakage or spontaneous/stimulated electronic emission), the system is said to be strongly coupled. Generally, the strong-coupling regime is characterized by the coherent energy exchange between the photon field and the electronic emitter (*i.e.* Rabi oscillations). In this regime, the electronic and photon subsystems can no longer be treated separately, making the accurate simulation of polariton dynamics very challenging. In this section, we explore various theoretical approaches in modeling the light-matter interactions present in optical cavities, such as Fabry-Pérot, where the coherent exchange in energy between all DOFs directly affects the resulting chemistry. Here, the electronic and photonic DOFs must be treated quantum mechanically on equal footing.

Recent experiments have shown a propensity to change chemistry via the coupling of quantized radiation and various molecular DOFs, namely electronic and vibrational strong coupling. The Fabry-Pérot-like cavities offer an extremely tunable cavity frequency (via the effective length between the cavity mirrors) while exhibiting widely varying coupling strengths highly dependent on the experimental setup<sup>53,147,147–155</sup>. There are many open questions regarding these experiments, such as the collective effects (*i.e.*, many-molecule or many-mode effects), which are present in many of the recent works due to the complexity of performing single-molecule experiments.

## 4.2 Theoretical and Computational Challenges

There are conceptual and technical challenges in performing simulations of polaritonic systems. The first theoretical hurdle is that of the Hamiltonian itself. For the Pauli-Fierz Hamiltonian (Eq. ??), the light-matter interaction requires explicit knowledge of the molecular dipole operator  $\hat{D}$  in the working basis. It turns out that a significant simplification can be made if one neglects all the contributions from the dipole matrix except for the ground-to-excited transition dipole matrix elements. There are two primary reasons for making this approximation: (I) The entirety of this quantity is not usually printed by default when executing standard electronic structure software for electronically excited states; however, the usual information given from these calculations is the ground-to-excited transition dipole moment,  $D_{0J}$ , which, for example in linear response calculations (e.g., TD-HF or TD-DFT), is a trivial quantity to achieve since the result of such schemes is the ground-to-excited transition density.<sup>156</sup> (II) Historically, the field of quantum optics was narrowly focused on the light-matter interaction between single atoms and the quantized cavity field. In such cases, the diagonal elements of the dipole operator are zero by construction, and the electronic energy differences between excited states were large, leading to neglecting of the excited-to-excited dipole coupling. With these approximations, one would find that a reduced Hamiltonian can be achieved, commonly referred to as the Jaynes-Cumming (JC) Hamiltonian.

## 4.3 Common Approximations Toward Historic Quantum Optics Models

The PF Hamiltonian (Eq. 11) is derived by applying the dipole approximation, which assumes the wavelength of the cavity fields is substantially larger than the matter system so that the spatial dependence of the transverse fields is neglected. In addition, many of the recent works on the simulation of *ab initio* polaritons have relied on approximate versions of the PF Hamiltonian that stem from historical applications in the quantum optics community.<sup>157–163</sup> In these heavily approximated Hamiltonians, usually a large truncation of the electronic and photonic subspaces is also performed such that only the ground and a single electronic excited state,  $|g\rangle$  and  $|e\rangle$  are included while only including the vacuum and singly excited Fock states,  $|0\rangle$  and  $|1\rangle$  in a single-mode cavity  $\alpha = 0$ . The total basis for this simple model is then confined to  $\{|g,0\rangle, |g,1\rangle, |e,0\rangle, |e,1\rangle\}$ . Note here that another approximation is that there is no permanent dipole in the ground  $|g\rangle$  or excited  $|e\rangle$  electronic states, i.e.,  $\mathbf{D}_{gg} = \mathbf{D}_{ee} = 0$ . The most commonly used Hamiltonian for modeling *ab initio* polaritons is the Jaynes-Cummings Hamiltonian  $\hat{H}_{JC}$ , which can be written as,

$$\hat{H}_{JC} = \hat{H}_m + \hat{H}_p + \sqrt{\frac{\omega_c}{2}} \boldsymbol{\lambda} \cdot \mathbf{D}_{ge} (\hat{\sigma} \hat{a}^\dagger + \hat{\sigma}^\dagger \hat{a}), \quad (43)$$

where  $\hat{\sigma}^\dagger$  ( $\hat{\sigma}$ ) is the creation (annihilation) operator for the molecule excitation between the ground  $g$  and excited  $e$  states. Here, two approximations have been made: (I) the rotating wave approximation (RWA) – which is to say, neglecting the highly oscillatory  $\hat{\sigma}^\dagger \hat{a}_\alpha^\dagger$  and  $\hat{\sigma} \hat{a}_\alpha$  terms in the light-matter interaction –

and (II) neglecting the DSE  $\hat{H}_{DSE}$ . This Hamiltonian is valid at ultra-low coupling strengths where the splitting between the one-photon-dressed ground state  $|g,1\rangle$  and the excited state with zero photons  $|e,0\rangle$  exhibit linear splitting (e.g. Rabi splitting) with an increase in the coupling strength  $\boldsymbol{\lambda}$ . The other two basis states  $|g,0\rangle$  and  $|e,1\rangle$  are completely decoupled from the interaction.

Two other common approximations to the PF Hamiltonian are the Rabi  $\hat{H}_{Rabi}$  and the rotating wave approximation  $\hat{H}_{RWA}$ . These two approximate Hamiltonians can be written as,

$$\hat{H}_{Rabi} = \hat{H}_M + \hat{H}_P + \sqrt{\frac{\omega}{2}} \boldsymbol{\lambda} \cdot \hat{\mathbf{D}} (\hat{\sigma} + \hat{\sigma}^\dagger) (\hat{a}^\dagger + \hat{a}), \quad (44)$$

$$\hat{H}_{RWA} = \hat{H}_M + \hat{H}_P + \sqrt{\frac{\omega}{2}} \boldsymbol{\lambda} \cdot \mathbf{D}_{ge} (\hat{\sigma}_{ge} \hat{a}^\dagger + \hat{\sigma}_{ge}^\dagger \hat{a}) + \hat{H}_{DSE}, \quad (45)$$

where the Rabi Hamiltonian  $\hat{H}_{Rabi} = \hat{H}_{PF} - \hat{H}_{DSE}$  and the RWA Hamiltonian applies the RWA to the PF Hamiltonian. Here,  $\hat{\sigma}_{ge} = |g\rangle \langle e|$  is the annihilation operator of the two-level electronic system. A more in-depth discussion on these Hamiltonians and the effects of the dipole self-energy term can be found in Refs. 164, 157, 165, and 166.

Note that in a two-state electronic system with only  $g$  and  $e$  states with no permanent dipole – e.g., historically, a single-atom system – then the DSE term only provides a uniform energy shift to the overall energy, similar to the zero-point energy of the photon Hamiltonian. In this sense, neglecting this term historically was well-motivated, but in systems with permanent dipoles in the ground or excited states (e.g., Li-F) or a system with many electronic states (e.g., molecules or solid-state materials), then the DSE term provides non-trivial physics and must be included. More recently, a many-level, many-mode generalization of these approximated Hamiltonians has been used, which retains the main approximations in each but is now not identical to the historical definition of each.<sup>157</sup>

Since the correct PF Hamiltonian in the dipole gauge has been known, the question remains of why the community returns to the approximated Hamiltonians for *ab initio* as well as model calculations. There are many subtleties to using the full PF Hamiltonian. From the electronic structure perspective, the many-level dipole operator needs to be computed as well as its square. The DSE term provides a very complicated description of the system for large coupling strengths since the dipole matrix,  $\hat{\mathbf{D}}$ , in realistic molecules is far from sparse with its square leading to further complications.<sup>164,167</sup> This allows for strong coupling between arbitrary states that is not trivial to know *a priori* based on chemical or physical intuition. Often, these Hamiltonians are parameterized based on cavity-free electronic structure calculations to obtain the energies and dipoles of the electronic adiabatic states (see more details in Sec. 4.4.2). In this case, the number of included electronic (as well as photonic) basis states should be treated as a convergence parameter. In this case, the DSE causes mixing between far-separated-in-energy electronic states, which leads to slow convergence in the basis set size for the matter DOFs,<sup>157</sup> and in general poor results compared to benchmarks, especially at large light-matter coupling strengths when using a small electronic basis.<sup>168–170</sup> This is precisely why many authors are devel-

oping self-consistent formulations to construct the low-energy polaritonic states without the need to calculate all the high-energy electronic states (see Sec. 4.4.2).<sup>167,168,170–175</sup>. Further, when considering the quantum dynamical propagation of polaritons in *ab initio* systems, additional nuclear gradients are required compared to cavity-free simulations where only the gradients of the adiabatic state energies are required to propagate the quantum dynamics, namely the nuclear gradients on the adiabatic dipole matrix and its square (*i.e.*,  $\nabla_R \hat{\mathbf{D}}$  and  $\nabla_R \hat{\mathbf{D}}^2$ ).<sup>158</sup>

#### 4.4 Theoretical modeling of polariton chemistry

##### 4.4.1 Cavity Born-Oppenheimer approximation

Within the Born-Oppenheimer (BO) approximation,<sup>176</sup> the total electronic-photonic-nuclear can be factorized as,

$$\Phi(\mathbf{r}, \mathbf{R}, \mathbf{q}_\alpha) = \chi(\mathbf{R}) \Psi(\mathbf{r}, \mathbf{q}_\alpha; \mathbf{R}), \quad (46)$$

where  $\chi(\mathbf{R})$  and  $\Psi(\mathbf{r}, \mathbf{q}_\alpha; \mathbf{R})$  are the nuclear and polaritonic (*i.e.*, electronic and photonic) wavefunctions, respectively. Note here that the polaritonic wavefunction is parameterized by the nuclear positions, exactly like the case without photonic DOFs outside the cavity. Further, one can invoke the usual Born-Huang-like expansion over the Born-Oppenheimer factorization as,

$$\Phi(\mathbf{r}, \mathbf{R}, \mathbf{q}_\alpha) = \sum_\mu \chi_a(\mathbf{R}) \psi_\mu(\mathbf{r}, \mathbf{q}_\alpha; \mathbf{R}), \quad (47)$$

where  $\psi_\mu(\mathbf{r}, \mathbf{q}_\alpha; \mathbf{R})$  are the BO wavefunctions analogous to those as outputted in standard electronic structure packages for the ground and excited adiabatic states. In this basis, which we will call the adiabatic polaritonic basis to draw a direct connection to the bare electronic case, we will discuss various ways to calculate such polaritonic wavefunctions  $\psi_\mu(\mathbf{r}, \mathbf{q}_\alpha; \mathbf{R})$  from *ab initio* calculations in a variety of approaches and levels of approximation.

In the following three sections, we will explore ways to obtain rigorous nuclear-position-parameterized wavefunctions for the entangled adiabatic electron-photon states  $\psi_\mu(\mathbf{r}, \mathbf{q}_\alpha; \mathbf{R})$ . First a brief description of a direct diagonalization approach (Sec. 4.4.2) with Hamiltonians parameterized with information from standard electronic structure, while the following sections (Secs. 4.4.3 and 4.4.4) will focus on the self-consistent approach toward re-developing the standard many-body schemes in electronic structure theory (*e.g.*, HF, DFT, CCSD, etc.) for the QED Hamiltonian (*e.g.*, QED-HF, QED-DFT, etc.).

In principle, one can keep a less restrictive expression for the total wavefunction which is a single exact product of the two distributions in three possible factorization schemes,

$$\begin{aligned} \Phi(\mathbf{r}, \mathbf{R}, \mathbf{q}_\alpha; t) &= \chi(\mathbf{R}; t) \Psi(\mathbf{r}, \mathbf{q}; \mathbf{R}, t) \\ &= \chi(\mathbf{r}; t) \Psi(\mathbf{R}, \mathbf{q}; \mathbf{r}, t) \\ &= \chi(\mathbf{q}; t) \Psi(\mathbf{r}, \mathbf{R}; \mathbf{q}, t) \end{aligned} \quad (48)$$

where  $\chi$  is the marginal probability distribution chosen to be the exact distribution for either of the three unique DOFs. The other factor  $\Psi$  describes the combined wavefunction of the left-

over DOFs that, for example, when  $\Psi = \Psi(\mathbf{r}, \mathbf{q}; \mathbf{R}, t)$ , satisfies,

$$1 = \prod_\alpha \int d\mathbf{q}_\alpha d\mathbf{r} \Psi(\mathbf{r}, \mathbf{q}_\alpha; \mathbf{R}, t). \quad (49)$$

In this formalism, the exact time-dependent dynamics is solved via a set of coupled equation that involves the time-dependent potential energy surface (TDPES)  $\epsilon(\mathbf{R}, t) = \langle \Psi(\mathbf{r}; \mathbf{R}, t) | \hat{H}_{\text{el}}(\mathbf{r}; \mathbf{R}, t) - i\partial_t | \Psi(\mathbf{r}; \mathbf{R}, t) \rangle$  dictated by the instantaneous  $\Psi$  at  $\mathbf{R}$  and the time-dependent vector potential which takes the form  $A(\mathbf{R}; t) = \langle \Psi(\mathbf{r}; \mathbf{R}, t) | -i\nabla | \Psi(\mathbf{r}; \mathbf{R}, t) \rangle$ . More details can be found in the original works.<sup>177–185</sup>

##### 4.4.2 Direct Diagonalization

Historically, the application of QED theory was focused on atomic physics. Here, single atoms were subjected to a cavity photon yielding Jaynes-Cummings-like linear splitting (see Eq. 43) since the atomic dipole matrix was approximated as a single matrix element between the ground and excited state. In this approximation, the QED Hamiltonian was “parameterized” by the energies of two electronic adiabatic states,  $E_g, E_e$ , and transition dipole element  $\mu_{eg}$  connecting the ground and excited state. For *ab initio* systems, this single dipole matrix element can be approximated by a Taylor-Series expansion around the Frank-Condon point.<sup>159,161,163</sup> In this way, one can directly diagonalize this 4x4 Hamiltonian in the  $\{|g, 0\rangle, |g, 1\rangle, |e, 0\rangle, |e, 1\rangle\}$  using standard, exact diagonalization techniques. The left basis  $\{|g\rangle, |e\rangle\}$  states are the electronic adiabatic states that are eigenfunctions of the electronic Hamiltonian  $\hat{H}_{\text{el}} = \hat{T}_e + \hat{V} = \hat{H}_M - \hat{T}_R$ , where  $\hat{V}$  includes all Coulomb interactions between nuclei and electrons. The right basis states  $\{|0\rangle, |1\rangle\}$  are the vacuum and single-photon Fock states, which are eigenstates of the photonic Hamiltonian  $\hat{H}_p = \hbar\omega_\alpha(\hat{a}_\alpha^\dagger \hat{a}_\alpha + \frac{1}{2})$ . In this minimal basis, the JC Hamiltonian can be written as,

$$(\hat{H}_{\text{JC}})_{IJ,nm} \doteq \begin{bmatrix} E_g & 0 & 0 & 0 \\ 0 & E_e & \sqrt{\frac{\omega_c}{2}} \boldsymbol{\lambda} \cdot \mathbf{D}_{eg} & 0 \\ 0 & \sqrt{\frac{\omega_c}{2}} \boldsymbol{\lambda} \cdot \mathbf{D}_{eg} & E_g + \hbar\omega_0 & 0 \\ 0 & 0 & 0 & E_e + \hbar\omega_0 \end{bmatrix}, \quad (50)$$

where the lowest and highest-energy basis states  $\{|g, 0\rangle, |e, 1\rangle\}$  are completely decoupled from the other basis states as well as each other. In this sense, the polaritonic ground state  $|\Psi_{\mu=0}\rangle = |g, 0\rangle$  and the fourth polaritonic state  $|\Psi_{\mu=4}\rangle = |e, 1\rangle$ , which is valid for ultra-low light-matter coupling strengths where the linear splitting is valid at the resonance condition. For non-linear effects, such as those including permanent dipoles, off-resonant excitations, or simply the exploration of the ground polaritonic state. To make this leap, one needs to make two major changes: (I) invoke the more rigorous Pauli-Fierz (PF) Hamiltonian  $\hat{H}_{\text{PF}}$  (see Eq. 11) and (II) treat the number of basis electronic and photonic states convergence parameters.

The matrix elements of the single-mode PF Hamiltonian in this same adiabatic-Fock basis – but now extended to include, in principle, an infinite number of electronic and photonic states – can

be written as,

$$\begin{aligned} (\hat{H}_{\text{PF}})_{IJ,nm} &= \left[ E_I + \omega_\alpha (n + \frac{1}{2}) \right] \delta_{IJ} \delta_{nm} \\ &+ \sqrt{\frac{\omega_\alpha}{2}} \boldsymbol{\lambda} \cdot \mathbf{D}_{IJ} (\sqrt{m+1} \delta_{n,m+1} + \sqrt{m} \delta_{n,m-1}) \\ &+ \frac{1}{2} \sum_K^{\mathcal{N}_{\text{el}}} (\boldsymbol{\lambda} \cdot \mathbf{D}_{IK})(\boldsymbol{\lambda} \cdot \mathbf{D}_{KJ}) \delta_{nm}, \end{aligned} \quad (51)$$

where  $\mathcal{N}_{\text{el}}$  is the number of adiabatic electronic states included in the basis. It is important to note that in this basis, the PF Hamiltonian is extremely sparse since the coupling elements only connect adjacent Fock states via the molecular dipole matrix since the matrix elements of the photonic coordinate  $\hat{q}$  are that of the harmonic oscillator (*i.e.*, only have nonzero super- and sub-diagonal elements). Further, the DSE contributions, in general, connect all of the electronic states of the system (with the same photon number) and is the most non-trivial aspect of this Hamiltonian and will vary strongly between molecular systems.

In this section, we have chosen to expand on the JC and PF Hamiltonians, confining ourselves to the adiabatic-Fock basis; however, other options exist, such as the generalized coherent states<sup>172,186,187</sup> or polarized Fock states.<sup>188</sup> In both of these bases, the photonic states are chosen such that the molecular dipole parameterizes the photonic state, thereby, in principle, reducing the convergence of the photonic basis. Further, this “direct diagonalization” approach to solving the PF Hamiltonian is dependent on this basis convergence, and the electronic basis is much more rigid since the adiabatic basis is ubiquitously used for its convenience. However, the convergence of this basis has yet to be thoroughly tested for a wide range of systems in solving the PF Hamiltonian, but it is expected to converge slowly due to the contributions from the DSE term.<sup>157,167</sup> This evidences the need to move to a more rigorous self-consistent solution for the polaritonic adiabatic states as is done for the electronic adiabatic states themselves.

#### 4.4.3 Self-consistent Polaritonic Single-particle Approaches

##### Ground State scQED Hartree-Fock

The mean-field approach to the QED Hamiltonian can be first cast in an identical way as the standard Hartree-Fock procedure in a larger Hilbert space, including the photonic states. A useful basis, referred to as generalized coherent states (GCS), can be performed such that the light-matter interaction part of the PF Hamiltonian can be shifted away. In the following discussion, we use the notation of Ref. 172; although many works use a similar scheme.<sup>172,187,189</sup> Assuming that the ground state HF wavefunction is  $|\text{HF}\rangle$  without the influence of the cavity, then the PF Hamiltonian can be partially diagonalized in the electronic subspace as,

$$\begin{aligned} \langle \text{HF} | \hat{H}_{\text{PF}} | \text{HF} \rangle &= E_{\text{HF}} + \omega_\alpha (\hat{a}_\alpha^\dagger \hat{a}_\alpha + \frac{1}{2}) \\ &+ \sqrt{\frac{\omega_\alpha}{2}} \langle \boldsymbol{\lambda}_\alpha \cdot \hat{\mathbf{D}} \rangle_{\text{HF}} \cdot (\hat{a}_\alpha^\dagger + \hat{a}_\alpha) + \frac{1}{2} \langle (\boldsymbol{\lambda}_\alpha \cdot \hat{\mathbf{D}})^2 \rangle_{\text{HF}}, \end{aligned} \quad (52)$$

where  $\langle \cdots \rangle_{\text{HF}} = \langle \text{HF} | \cdots | \text{HF} \rangle$  is the HF ground state expectation value of the electronic subsystem. Introducing the GSC transformation as,

$$\hat{U}(z_\alpha) = e^{z_\alpha \hat{a}_\alpha^\dagger - z_\alpha^* \hat{a}_\alpha}, \quad (53)$$

where  $\mathbf{z} = (z_0, z_1, \dots)$  is a vector of arbitrary complex numbers specific to each cavity mode, we can completely remove the light-matter interaction from the Hamiltonian with a specific choice of  $z_\alpha$  as,

$$\mathbf{z} \rightarrow -\frac{\langle \boldsymbol{\lambda}_\alpha \cdot \hat{\mathbf{D}} \rangle_{\text{HF}}}{\sqrt{2\omega_\alpha}}. \quad (54)$$

This rotation results in a shift of the photonic operators by  $\mathbf{z}$ . Transforming  $\hat{H}_{\text{PF}}$  as  $\hat{U}(\mathbf{z}) \langle \text{HF} | \hat{H}_{\text{PF}} | \text{HF} \rangle \hat{U}^\dagger(\mathbf{z}) = \hat{H}_{\text{pl}}(\mathbf{z})$ , which can be written as,

$$\begin{aligned} \hat{H}_{\text{pl}}(\mathbf{z}) &= \langle \text{HF} | \hat{U}(\mathbf{z}) \hat{H}_{\text{PF}} \hat{U}^\dagger(\mathbf{z}) | \text{HF} \rangle \\ &= E_{\text{HF}} + \omega_\alpha (\hat{a}_\alpha^\dagger \hat{a}_\alpha + \frac{1}{2}) + \frac{1}{2} \langle (\boldsymbol{\lambda}_\alpha \cdot \Delta \hat{\mathbf{D}})^2 \rangle_{\text{HF}}, \end{aligned} \quad (55)$$

where the light-matter coupling is inherently removed in this basis since  $\langle \Delta \hat{\mathbf{D}} \rangle_{\text{HF}} = \langle \hat{\mathbf{D}} - \langle \hat{\mathbf{D}} \rangle_{\text{HF}} \rangle_{\text{HF}} = 0$ . Note that this removal only exists for mean-field-like methods, which do not include direct interactions between the ground and excited states. Now, the light-matter interaction only appears through the DSE-like term, which now depends not on the dipole  $\mathbf{D}$  but rather on the variance in the ground state permanent dipole,

$$\langle (\Delta \hat{\mathbf{D}})^2 \rangle_{\text{HF}} = \langle \hat{\mathbf{D}}^2 \rangle_{\text{HF}} - \langle \hat{\mathbf{D}} \rangle_{\text{HF}}^2, \quad (56)$$

which can be understood as dipole fluctuations inducing light-matter interactions. In this basis, the Hamiltonian can now be diagonalized in the photonic sub-space with photon number states. The ground polaritonic state is then defined as the product  $|\Psi_0(\mathbf{r}, \mathbf{q}_\alpha; R)\rangle = |\text{HF}\rangle \otimes |0\rangle$ . The HF energy can be written as,

$$\begin{aligned} E_{\text{QED-HF}} &= E_{\text{HF}} + \sum_\alpha \frac{1}{2} \langle (\boldsymbol{\lambda}_\alpha \cdot \Delta \hat{\mathbf{D}})^2 \rangle_{\text{HF}} \\ &= E_{\text{HF}} + \sum_{iv} \sum_\alpha (\boldsymbol{\lambda}_\alpha \cdot \mathbf{D}_{ia})^2 \end{aligned} \quad (57)$$

where  $i$  and  $a$  are the occupied and virtual single-particle state labels. This energy is variationally minimized to achieve convergence in the basis of canonical spin-orbitals.

##### QED Density Functional Theory

Similarly to the scQED-HF approach, the self-consistent QED density functional theory (scQED-DFT) is composed in a similar way, where the main elements of DFT remain, such as the exchange-correlation functional of the density. However, there are additional photonic DOF that influence the electronic subsystem. As for the scQED-HF case, the generalized coherent state basis removes the light-matter coupling term while leaving the DSE term. In this sense, the energy of the Kohn-Sham energy can be written as,

$$E_{\text{QED-DFT}} = E_{\text{DFT}} + \frac{1}{2} \sum_\alpha \langle (\boldsymbol{\lambda}_\alpha \cdot \hat{\mathbf{D}})^2 \rangle, \quad (58)$$

which is then minimized with respect to the Kohn-Sham spin-

orbitals. There are many ways to set up the scQED-DFT problem, such as employing a novel exchange-correlation functional to account for light-matter correlation effects<sup>190–193</sup> or working with electron-only exchange-correlation functionals using the coherent state basis for the photonic DOFs.<sup>157,171,172</sup> In any case, the resulting ground state is uncorrelated by nature of the DFT formalism.

### Excited States scQED-TD-(HF,DFT)

The time-dependent analogues to the aforementioned single-particle approaches are powerful tools to probe non-equilibrium densities that give rise to electronic excited states. One of the most popular approaches is one of linear response (LR), resulting in LR-TD-HF and LR-TD-DFT for bare molecular systems in the random phase approximation (RPA). Although, it should be noted that the real-time propagation of the single-particle density matrix – leading to the real-time TD-HF and real-time TD-DFT approaches – is, in principle, a more robust approach but one that is usually more costly than that of linear response. Such schemes have already been developed for the simulation of molecular polaritons using classical photon DOFs.<sup>194,195</sup> Here, we will focus our attention on the LR formalism, specifically using a Casida-like approach to writing the random phase approximation (RPA), originally formulated by Flick and co-workers<sup>168</sup> and rewritten in the language of Casida by the groups of Shao<sup>157</sup> and De-Prince.<sup>171</sup> It should be noted that other formulations of CIS-like excited states can be found in the community, such as the non-Hermitian CIS aimed at simulating cavity loss via a complex photon frequency<sup>170</sup>.

As per usual, and following the notation of Ref. 171, the LR-TD-HF and LR-TD-HF eigenvalue equations using the Casida formalism can be written to include the QED components that satisfy the Pauli-Fierz Hamiltonian (Eq. 11) as,

$$\begin{bmatrix} \mathbf{A} + \Delta & \mathbf{B} + \Delta' & \mathbf{g}^\dagger & \mathbf{g}^\dagger \\ \mathbf{B} + \Delta' & \mathbf{A} + \Delta & \mathbf{g}^\dagger & \mathbf{g}^\dagger \\ \mathbf{g} & \mathbf{g} & \omega & \mathbf{0} \\ \mathbf{g} & \mathbf{g} & \mathbf{0} & \omega \end{bmatrix} \begin{bmatrix} \mathbf{X} \\ \mathbf{Y} \\ \mathbf{N} \\ \mathbf{M} \end{bmatrix} = \Omega \begin{bmatrix} \mathbf{1} & \mathbf{0} & \mathbf{0} & \mathbf{0} \\ \mathbf{0} & -\mathbf{1} & \mathbf{0} & \mathbf{0} \\ \mathbf{0} & \mathbf{0} & \mathbf{1} & \mathbf{0} \\ \mathbf{0} & \mathbf{0} & \mathbf{0} & -\mathbf{1} \end{bmatrix} \begin{bmatrix} \mathbf{X} \\ \mathbf{Y} \\ \mathbf{N} \\ \mathbf{M} \end{bmatrix} \quad (59)$$

where  $\mathbf{A}$  and  $\mathbf{B}$  are the usual excitation and de-excitation block matrices written using the TD-HF and TD-DFT approximations,

$$A_{ia,jb}^{\text{TD-HF}} = \delta_{ij}\delta_{ab}(\epsilon_i - \epsilon_a) + \langle ja||bi\rangle, \quad B_{ia,jb}^{\text{TD-HF}} = \langle ij||ab\rangle \quad (60)$$

$$A_{ia,jb}^{\text{TD-DFT}} = \delta_{ij}\delta_{ab}(\epsilon_j - \epsilon_a) + f_{ja,bi}^{\text{h/xc}}, \quad B_{ia,jb}^{\text{TD-DFT}} = f_{ij,ab}^{\text{h/xc}} \quad (61)$$

with  $\langle ja||bi\rangle = \langle ja|bi\rangle - \langle ja|ib\rangle$  as the standard two-body interaction terms,  $f_{ia,jb}^{\text{h/xc}}$  as the combination of the Hartree (h) and exchange-correlation (xc) terms from a standard DFT formalism, and  $\epsilon_a$  is the energy of single-particle orbital  $a$ . Recall that  $i, j$  denote occupied orbitals while  $a, b$  denote unoccupied orbitals. The

QED-specific terms can be written similarly as,

$$g_{ia} = \sum_{\alpha} \sqrt{\frac{\omega_{\alpha}}{2}} D_{ia}^{\lambda_{\alpha}} \quad (62)$$

$$\Delta_{ia,jb} = \sum_{\alpha} D_{ia}^{\lambda_{\alpha}} D_{bj}^{\lambda_{\alpha}} - D_{ij}^{\lambda_{\alpha}} D_{ab}^{\lambda_{\alpha}} \quad (63)$$

$$\Delta'_{ia,jb} = \sum_{\alpha} D_{ia}^{\hat{u}_{\alpha}} D_{jb}^{\lambda_{\alpha}} - D_{ib}^{\lambda_{\alpha}} D_{aj}^{\lambda_{\alpha}} \quad (64)$$

Here,  $D_{ia}^{\lambda_{\alpha}} = \lambda_{\alpha} \cdot \mathbf{D}_{ia}$  with  $\hat{u}_{\alpha}$  as the polarization of the  $\alpha$ <sub>th</sub> cavity mode. Further,  $\Delta = \Delta'$  if the molecular orbitals are real-valued.<sup>171</sup>

An important distinction between various implementations of the QED-TD-DFT approaches in the community is whether the single-particle orbitals used in the formulation are “relaxed” in the presence of the cavity or are simply the bare electronic single-particle states. For example, in Ref. 157, the orbitals are not relaxed while in the Refs. 168 and 171 the orbitals are relaxed. While it is clear that using a relaxed reference state for the basis of the RPA equations would provide a more rigorous result, it is not clear whether identical results can be obtained in the infinite basis limit of both approaches, *i.e.*, including more single-particle states in the CIS-like expansion in excited Slater determinants. Since the RPA equations are iteratively solved, the expansion coefficients of the excited Slater determinants may result in the same excited state observables in the infinite basis limit, while for a finite basis, it may not.

#### 4.4.4 Self-consistent Polaritonic Coupled Cluster Approaches (CC,EOM-CC)

Despite computational efficiency, DFT or mean-field HF methods usually underestimate the correlations. In particular, the mean-field method cannot describe the electron-photon correlation, and the exchange-correlation function for electron-photon interaction is unknown. Consequently, the QED counterpart of coupled-cluster theory (QED-CC) is proposed.<sup>36,37,173–175,196,197</sup> Similar to the conventional CC theory, QED-CC employs an exponential wavefunction Ansatz to derive the ground state,

$$|\Psi_{CC}\rangle = e^{\hat{T}} |\Phi_0\rangle, \quad (65)$$

where  $|\Phi_0\rangle$  is the reference wave function which is usually chosen to be the tensor product of Hartree-Fock (HF) determinant and photon vacuum state, *i.e.*,  $|\Phi_0\rangle = |\phi_0\rangle \otimes |0\rangle$ .  $\hat{T}$  is defined in excitation configurations. Within QED-CC theory, the generalized excitation operator includes three components,  $\hat{T} = \hat{T}_e + \hat{T}_p + \hat{T}_{ep}$ , including electronic ( $\hat{T}_e$ ), photonic ( $\hat{T}_p$ ), and coupled electronic-

photonic ( $\hat{T}_{ep}$ ) excitations,

$$T_e = \sum_{ia} t_i^a \hat{a}_a^\dagger \hat{a}_i + \sum_{ijab} t_{ij}^{ab} \hat{a}_a^\dagger \hat{a}_b^\dagger \hat{a}_j \hat{a}_i + \dots \equiv \sum_\mu^{N_e} t_\mu \hat{\tau}_\mu, \quad (66)$$

$$\hat{T}_p = \sum_\alpha \gamma_\alpha \hat{b}_\alpha^\dagger + \frac{1}{2} \sum_{\alpha\beta} \gamma_{\alpha\beta} \hat{b}_\alpha^\dagger \hat{b}_\beta^\dagger + \dots \equiv \sum_n^{N_p} \gamma_n \hat{B}_n, \quad (67)$$

$$\hat{T}_{ep} = \sum_{ia,\alpha} t_i^a \hat{a}_a^\dagger \hat{a}_i \hat{b}_\alpha^\dagger + \frac{1}{2} \sum_{ijab,\alpha\beta} t_{ij}^{ab} \hat{a}_a^\dagger \hat{a}_b^\dagger \hat{a}_j \hat{a}_i \hat{b}_\alpha^\dagger \hat{b}_\beta^\dagger + \dots \equiv \sum_{\mu,n} \chi_{\mu,n} \hat{\tau}_\mu \hat{B}_n. \quad (68)$$

where  $N_e$  and  $N_f$  are the numbers of electrons and photon Fock states, respectively.  $\hat{\tau}_\mu = \prod_k^\mu \hat{E}_{ik}^{ak}$  and  $\hat{E}_i^a = \hat{a}_a^\dagger \hat{a}_i + h.c.$  are the  $\mu$ -body and single-body excitation operators, respectively.  $\hat{B}_n = \prod_\alpha^n \hat{b}_\alpha^\dagger$  is the  $n$ -body photonic excitation. The parameters  $(t_\mu, \gamma_n, \chi_{\mu,n})$  are the cluster amplitudes, which are determined through a set of CC equations. The cluster operator  $\mu \in \{\hat{\tau}_\mu, \hat{B}_n, \hat{\tau}_\mu \hat{B}_n\}$  generates a set of orthogonal excited configurations  $(\{|\mu\rangle\})$ ,

$$|\mu\rangle = \hat{\mu} |\Phi_0\rangle. \quad (69)$$

The combined set  $(\{|\Phi_0\rangle, |\mu\rangle\})$  defines the Hilbert subspace for both ground and excited states. Using the CC ansatz in Eq. 65, the time-dependent Schrödinger equation leads to

$$\hat{H}_{PF} |\Psi_{CC}\rangle = E_{cc} |\Psi_{CC}\rangle, \quad (70)$$

A set of Slater determinants  $(|HF\rangle, |\mu\rangle)$  is used as the basis set to solve the Eq. 70, which leads to

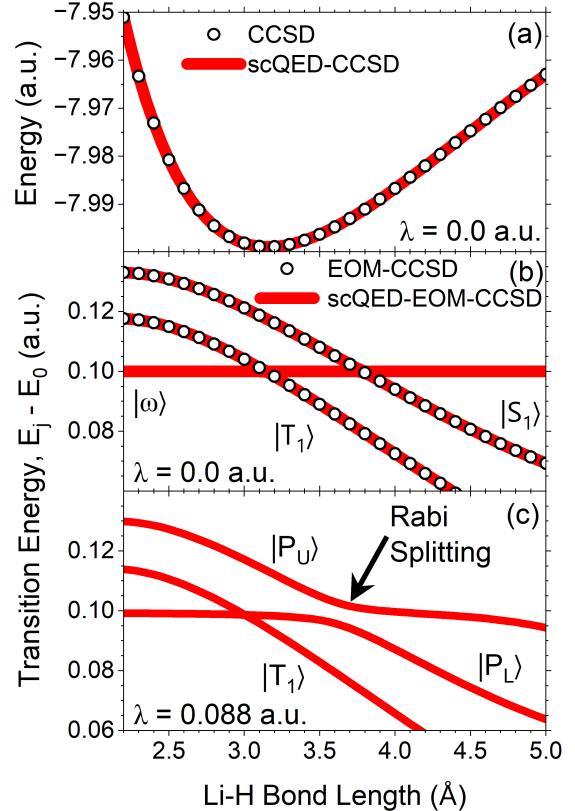
$$E_{CC} = \langle HF | \hat{H}_{PF} | HF \rangle. \quad (71)$$

Where  $\hat{H}_{PF} = e^{-\hat{T}} \hat{H}_{PF} e^{\hat{T}}$  is the dressed Hamiltonian. The cluster amplitudes are determined from the projected equation,

$$\Omega_\mu \equiv \langle \mu | e^{-\hat{T}} \hat{H} e^{\hat{T}} | \Phi_0 \rangle = 0 \quad (72)$$

The scQED-CC theory leads to the FCI solution if no truncation on the excitation operator  $\hat{T}$  is applied. However, in order to trade-off between accuracy and computational efficiency, the excitation operator is usually truncated at the doubles (CCSD) level. Some tests have been performed on the level of truncation in the photonic excitations (up to 10) in the CC operator for model systems.<sup>196</sup> However, the scaling of scQED-CCSD is  $\mathcal{O}(N_{el}^6 N_p^{M_p})$  in general, where  $M_p$  is the number of photon modes. Such a scaling makes it usually a bit expensive to rigorously test in extended molecular systems that include many electrons and/or when many photon modes and Fock states are used in the calculations. To further reduce the computational cost, the generalized coherence state<sup>198</sup> may be used to reduce the photon basis in the scQED-CC calculations.

It should be noted that CC methods exhibit size extensivity even in its truncated forms, meaning that the sum of the energies of non-interacting subsystems is equal to the total energy. This is in contrast to CI approaches, where errors in extensivity can become increasingly large with an increasing number of subsystems. In addition, CC theory has the advantage of size extensivity in that



**Fig. 5** Polariton states of LiH as a function of Li-H bond length. (a) CCSD (black circles) and scQED-CCSD (solid red curve) ground states at zero light-matter coupling strength  $\lambda = 0$  a.u. (b) EOM-CCSD (black circles) triplet  $|T_1\rangle$  and singlet  $|S_1\rangle$  states as well as the low-lying states  $(|T_1\rangle, |\omega\rangle, \text{ and } |S_1\rangle)$  calculated at scQED-EOM-CCSD with light-matter coupling strength  $\lambda = 0$  a.u. The plot clearly shows the photon line ( $|\omega\rangle$ ) and perfect overlap of the  $S_1$  states. (c) Upper ( $|P_U\rangle$ ) and lower ( $|P_L\rangle$ ) polariton states with  $\lambda = 0.088$  a.u. with the triplet state  $|T_1\rangle$  uncoupled.

the excitation energies of each subsystem do not vary with the size of the total system when the subsystems are non-interacting.

In addition, the excited polariton states can be computed with the corresponding EOM formalism, which parameterizes a neutral or charged excitation by applying an excitation operator to the CC ground state<sup>199</sup>

$$|\hat{R}\rangle = \hat{R} |\Phi_{cc}\rangle = \hat{R} e^{\hat{T}} |\Phi_0, 0\rangle. \quad (73)$$

Where  $\hat{R}$  also include the electronic ( $\hat{R}_e$ ), photonic ( $\hat{R}_p$ ), and coupled electronic-photonic ( $\hat{R}_{ep}$ ) excitations. Because the excitation operator,  $\hat{R}$ , commutes with the excitation operators in  $\hat{T}$ , solving this eigenvalue problem is equivalent to finding a right eigenvector of the similarity transformed Hamiltonian,

$$\langle \mu | \hat{H} \hat{R}^n | \Phi_0, 0 \rangle = E_n \hat{R}_\mu^n. \quad (74)$$

Here,  $E_n$  is the energy of the nth excited state, and  $\mu$  indexes an element of the excitation operator  $\hat{R}$ . The excitation operator,  $\hat{R}$ , can be chosen to access charged or neutral excitations.

In practice, the eigenvalue problem is solved by iterative diag-

onalization. The necessary equations for the sigma vectors are derived and implemented efficiently.

The sigma equations of polaritonic EOM-CCSD are defined as<sup>199</sup>

$$\begin{aligned}\sigma_{m\mu} &= \langle \mu, m | \bar{H} \hat{R} | \Phi_0, 0 \rangle \\ &= \langle \Phi_0, 0 | \tau_\mu^\dagger(\hat{b})^m \bar{H} \hat{R} | \Phi_0, 0 \rangle.\end{aligned}\quad (75)$$

(note,  $|\mu, m\rangle = \tau_\mu(\hat{b}^\dagger)^m |\Phi_0, 0\rangle$  and  $\tau_\mu$  is single or double excitation operator. While this approach provides the correct equations, the resulting equations contain hundreds of terms, even for simple EOM theories, and are, therefore, difficult to optimize. Hence, Garnet's paper suggested separately generating the different sectors of the similarity transformed Hamiltonian,

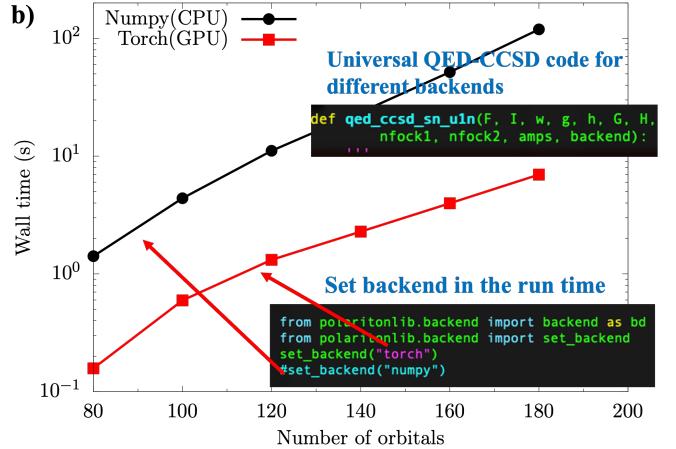
$$\bar{H}_{m\mu,n\nu} = \langle \Phi_0, 0 | \tau_{\mu,m}^\dagger \bar{H} \tau_{n,\nu} | \Phi_0, 0 \rangle. \quad (76)$$

for all relevant excitations. Then generate equations for the different sectors of the configuration-interaction-like sigma vector as

$$\sigma_{m\mu} = \sum_{n\nu} \bar{H}_{m\mu,n\nu} R_{n\nu}. \quad (77)$$

We recently implemented such QED-CCSD/EOM-CCSD method to compute the polariton states. With the diagrammatic technique, an auto code generator and optimizer are developed to generate the QED-CCSD/EOM-CCSD equations to arbitrary photon order. Fig. 5 is an example of the Li-H bond dissociation curves in a single-mode optical cavity. Fig. 5a shows the ground state Born-Oppenheimer potential energy surface for the bare electronic system (black circles) and for the polaritonic system (red curve) at zero light-matter coupling strength ( $\lambda = 0$ ). The excited states inside (red curves) and outside (black circles) of the cavity are shown at zero coupling strength ( $\lambda = 0$ ), where two multiplicities (singlet  $|S_1\rangle \equiv |S_1, 0\rangle$  and triplet  $|T_1\rangle \equiv |T_1, 0\rangle$ ) of electronic states are shown as well as a single cavity state ( $|\omega\rangle \equiv |S_0, 1\rangle$ ). Here, we have used the notation  $|S_J\rangle \otimes |n\rangle = |S_J, n\rangle$  where the left ket in the product signifies the electronic DOFs while the right signifies the photonic Fock state label (*i.e.*, the number of photons in the photon-dressed electronic state). At finite light-matter coupling of  $\lambda = 0.088$  a.u., the cavity state with one photon  $|\omega\rangle$  couples strongly with the singlet excited electronic state with zero photons  $|S_1\rangle$  while the excited electronic triplet state  $|T_1\rangle$  is negligibly affected. The so-called Rabi splitting appears at the degenerate point between the singlet electronic state and the cavity state,  $R_{\text{Li}-\text{H}} \approx 3.75$ , forming the upper ( $|P_U\rangle$ ) and lower ( $|P_L\rangle$ ) polaritonic states with mixed electronic and photonic character.

The computational efficiency of many-body electronic (or polaritonic) structure codes is nearly as important as the method itself. We have implemented various backend options for our code (presented in Fig. 5). The efficiency of Numpy *einsum* (CPU-accessible) and Torch *einsum* (GPU-accessible) functions are shown in Fig. 6 on a vertical log scale as a function of the number of orbitals (*i.e.*, electrons) included in the calculation. The GPU hardware allows the QED-CC code to consistently operate with an order of magnitude less wall time than the CPU



**Fig. 6** Example of running calculations with our modular QED-CC code on different architectures (CPU vs GPU on a desktop) by simply setting the backend (see the inset code).  $10\times - 20\times$  speed up is observed on Torch GPU.

version. These results indicate that the conversion toward GPU hardware is required for exploring the frontier science of molecular polaritons.

## 5 Perspective and summary

### 5.1 Multiscale method for light-matter interaction

Though there have been extensive developments in QED methods, most of the current implementations are based on the dipole approximation. However, the dipole approximation can fail in many cases. The nanoplasmonic cavities allow for light localization into deeply subwavelength dimensions, leading to effective mode volumes as small as a few nm<sup>3</sup> or even Å<sup>3</sup> (microcavities)<sup>23</sup>. Consequently, the size of molecules becomes comparable to the cavity volumes, and the widely used dipole approximation breaks down. Moreover, the cavity confines the photon field in a certain direction when coupling molecules with many modes inside a cavity (either nanophotonic or nanoplasmonic ones). For a given probing angle  $\theta$  (relative to the normal vector of the cavity), the photon energy has a certain dispersion function. Under this situation, the dipole approximation along the in-plane direction of the cavity no longer holds as well<sup>159,200</sup>. Hence, a general method that can compute the multiple spatial-dependent cavity eigenmodes  $\lambda_{nk}(\mathbf{r})$  and light-matter couplings (from Maxwell's equations) is required.

As argued above, the EM field within the cavity is, in principle, not homogeneous, and spatial dependence matters in light-matter interactions. The spatial distribution of the EM field can be solved via standard computational electromagnetic methods, such as Finite-difference time-domain (FDTD)<sup>201</sup>, for Maxwell's equations. Consequently, the light-matter coupling strength can stem from a delicate balance between the spatial dependence of the electronic wavefunctions and the photonic fields. Therefore, only a quantum model fully incorporates the inhomogeneities of the exciton transition charge density can quantitatively describe this interplay. Using a fully first-principles methodology to de-

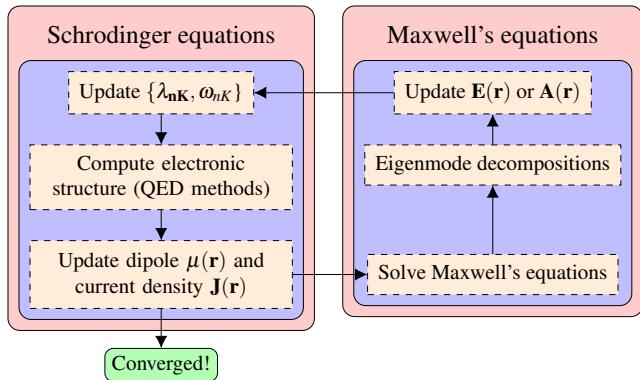
scribe the quantum chemistry of molecules placed inside the cavity, we can reveal the limitations of the point-dipole approach to address the exciton dynamics in weak and strong coupling regimes. Besides, current methods usually don't consider the feedback of molecular systems on the cavities. In many situations, modeling of molecular systems only is insufficient as the molecular response can significantly affect the EM distributions, especially in nanoplasmonic cavities. Even though the *ab initio* QED methods we previously developed take into account the interplay between the electronic and photonic DOFs, the molecular response on the EM environment is not considered.

In principle, we could solve the Schrödinger and Maxwell equations simultaneously in order to obtain access to the radiated fields and, with it to the self-consistent evolution of light and matter, i.e., the cavity photon modes can be affected by the modular dipoles due to the Ampere's Law (Helmholtz's equation):

$$\left[ \nabla \times \frac{1}{\mu_r(\mathbf{r}\omega)} \nabla \times -\omega^2 \mu_0 \epsilon_0 \epsilon(\mathbf{r}\omega) \right] \mathbf{E}(\mathbf{r}) = -e \mathbf{J}(\mathbf{r}), \text{ or} \quad (78)$$

$$\left( \nabla^2 - \frac{1}{c^2} \frac{\partial^2}{\partial t^2} \right) \mathbf{A} = -\mu_0 \mathbf{J}. \quad (79)$$

Where  $\mathbf{J}$  is the paramagnetic current density due to the molecular dipoles. And according to Maxwell's equations, each current induces an electromagnetic field for which its precise spatial and polarization structure depends on the electromagnetic environment—oscillating charges emit light. Hence, a fully self-consistent QED method should consider the feedback of molecular dipoles on the cavity properties. The flowchart is demonstrated in Figure 7.



**Fig. 7** Self-consistent loop between two sets of equations. The current density from the QED solvers is used to update EM fields. The self-consistency between Maxwell and Schrödinger equation can be bypassed (in the linear-response regime) by using the pre-calculated Dyadic Green's functions.

Alternatively, the generated field can be expressed with the help of the dyadic Green's tensor,

$$\mathbf{E}(\mathbf{r}, \omega) = i\mu_0 \omega \int d\mathbf{r}' G(\mathbf{r}, \mathbf{r}', \omega) [-e \mathbf{J}(\mathbf{r}', \omega)], \quad (80)$$

where Green's function is the formal solution of Helmholtz's equa-

tion

$$\left[ \nabla \times \frac{1}{\mu_r(\mathbf{r}\omega)} \nabla \times -\omega^2 \mu_0 \epsilon_0 \epsilon(\mathbf{r}\omega) \right] G(\mathbf{r}, \mathbf{r}', \omega) = \delta(\mathbf{r}, \mathbf{r}'), \quad (81)$$

with linear media  $\epsilon(\mathbf{r})$ . The Green's function in Eq. 81 can be solved from the Frequency domain FDTD methods.

Utilizing Dyadic Green's functions offers a powerful approach to simplify the description of certain components of a matter system while focusing on the computation of electronic structure only. Such a concept has been proposed in the past in a semi-classical treatment of light-matter interaction<sup>202,203</sup>. Introducing Green's function is particularly advantageous in multicomponent systems that span different length scales, such as a microscopic molecule (which is described by the current density  $\mathbf{J}$ ) and macroscopic solvent (represented by a parameterized dielectric function  $\epsilon(\mathbf{r})$ ). The EM environment embedded via Dyadic Green's function can be obtained numerically using many standard Maxwell solvers with certain boundary conditions, such as FDTD<sup>201</sup> and method of moments<sup>204</sup>, providing the photon mode structures that are coupled with QED electronic structure solvers. By computing  $\mathbf{G}$  beforehand, the self-consistency is embedded solely via the current density, eliminating the need to treat QED electronic structures and Maxwell's equation simultaneously. The bypassing of the self-consistency between Maxwell-Schrödinger equations is a significant strength of Dyadic Green's function approach. This makes the approach more accessible and easier to implement in various electronic structure solvers. But, *it should be noted that introducing linear media restricts the evaluation in the linear response regime*.

Moreover, the multiscale methods described above can have a significant influence other than polariton chemistry. It could affect the study of light-matter interactions in chemistry, physics, materials science, and energy science at large, such as plasmon chemistry<sup>21</sup>, quantum plasmonics<sup>205</sup>, quantum information transduction<sup>206</sup>, and photonic-microelectronics integration and polaritonic devices<sup>207</sup>. The integration with open quantum system method, particularly time-dependent quantum transport theories<sup>208–210</sup>, will also open the door to simulating light-driven quantum transport phenomena<sup>211,212</sup> in either weak or strong coupling regimes.

## 5.2 Summary

Controlling chemistry or molecular properties has been a long-standing Holy Grail for many decades. Only recently, new possibilities have emerged in the context of light-matter interactions, via either plasmon or polariton-mediated chemistry, which holds the promise of providing fundamental new strategies to control chemical reactions that are completely distinct from traditional ones (such as electrochemistry, thermochemistry, and photochemistry). Consequently, the application of light-matter interactions in manipulating chemistry has attracted increasing experimental and theoretical attention. However, the inherent multiscale nature of light-matter interaction problems poses significant challenges for both experimental and theoretical investigations of the underlying processes. Besides, multiscale processes at different

length and time scales make it difficult to optimize the performance of light-matter interaction-mediated chemistry, and precise modeling of these processes is essential for obtaining a comprehensive understanding of the fundamental mechanisms or current experiments. Hence, the development of multiscale theoretical and computational models that can precisely describe the dynamical processes in light-matter interaction-mediated chemistry across different time and length scales, in conjunction with massively parallel algorithms for large-scale simulations, is essential for obtaining comprehensive insights into the fundamental mechanisms of current experiments. These simulations allow for the modeling of complex, multiscale processes that would be otherwise difficult or impossible to observe experimentally alone. Such development will ultimately facilitate the discovery of new reaction pathways enhanced by light-matter interactions.

## Conflicts of interest

There are no conflicts to declare.

## Acknowledgements

The authors acknowledge support from the Laboratory Directed Research and Development (LDRD) program of Los Alamos National Laboratory.(Grant No. 20220527ECR). XL and YZ acknowledge support from the US DOE, Office of Science, Basic Energy Sciences, Chemical Sciences, Geosciences, and Biosciences Division under Triad National Security, LLC (“Triad”) contract Grant 89233218CNA000001 (FWP: LANLE3F2). LANL is operated by Triad National Security, LLC, for the National Nuclear Security Administration of the U.S. Department of Energy (contract no. 89233218CNA000001).

## Notes and references

- 1 G. R. Fleming and M. A. Ratner, *Phys. Today*, 2008, **61**, 28–33.
- 2 A. Frisk Kockum, A. Miranowicz, S. De Liberato, S. Savasta and F. Nori, *Nat. Rev. Phys.*, 2019, **1**, 19–40.
- 3 N. Rivera and I. Kaminer, *Nat. Rev. Phys.*, 2020, **2**, 538–561.
- 4 P. W. Anderson, *Science*, 1972, **177**, 393–396.
- 5 L. Novotny and B. Hecht, *Annual Review of Physical Chemistry*, 2006, **57**, 303–331.
- 6 J. Weiner and F. Nunes, *Light-Matter Interaction: Physics and Engineering at the Nanoscale*, Oxford University Press, 2012.
- 7 G. Mahan, *Many-Particle Physics*, Springer, 2000.
- 8 P. L. Stiles, J. A. Dieringer, N. C. Shah and R. P. Van Duyne, *Annu. Rev. Anal. Chem.*, 2008, **1**, 601–626.
- 9 S. Mukamel, *Annu. Rev. Phys. Chem.*, 2000, **51**, 691–729.
- 10 Y. Zhang, L. Meng, C. Yam and G. Chen, *J. Phys. Chem. Lett.*, 2014, **5**, 1272–1277.
- 11 C. Yam, L. Meng, Y. Zhang and G. Chen, *Chem. Soc. Rev.*, 2015, **44**, 1763–1776.
- 12 L. Meng, C. Yam, Y. Zhang, R. Wang and G. Chen, *J. Phys. Chem. Lett.*, 2015, **6**, 4410–4416.
- 13 L. Meng, Y. Zhang and C. Yam, *J. Phys. Chem. Lett.*, 2017, **8**, 571–575.
- 14 R. Wang, Y. Zhang, G. H. Chen and C. Y. Yam, *Prog. Electro-*
- magn. Res.*, 2015, **154**, 163–170.
- 15 C. L. Degen, F. Reinhard and P. Cappellaro, *Rev. Mod. Phys.*, 2017, **89**, 035002.
- 16 F. Flamini, N. Spagnolo and F. Sciarrino, *Rep. Prog. Phys.*, 2018, **82**, 016001.
- 17 L. Zhou, Y. Zhang, Z. Zhuo, A. J. Neukirch and S. Tretiak, *J. Phys. Chem. Lett.*, 2018, **9**, 6915–6920.
- 18 T. Mirkovic, E. E. Ostroumov, J. M. Anna, R. van Grondelle, Govindjee and G. D. Scholes, *Chem. Rev.*, 2017, **117**, 249–293.
- 19 P. Forn-Díaz, L. Lamata, E. Rico, J. Kono and E. Solano, *Rev. Mod. Phys.*, 2019, **91**, 025005.
- 20 M. L. Brongersma, N. J. Halas and P. Nordlander, *Nat. Nanotechnol.*, 2015, **10**, 25–34.
- 21 C. Zhan, J. Yi, S. Hu, X.-G. Zhang, D.-Y. Wu and Z.-Q. Tian, *Nat. Rev. Methods Primers*, 2023, **3**, 12.
- 22 E. Kazuma and Y. Kim, *Angew. Chem. Int. Ed.*, 2019, **58**, 4800–4808.
- 23 J. J. Baumberg, *Nano Lett.*, 2022, **22**, 5859–5865.
- 24 R. F. Ribeiro, L. A. Martínez-Martínez, M. Du, J. C. Gonzalez-Angulo and J. Yuen-Zhou, *Chem. Sci.*, 2018, **9**, 6325–6339.
- 25 T. W. Ebbesen, *Acc. Chem. Res.*, 2016, **49**, 2403–2412.
- 26 S. Kéna-Cohen and S. R. Forrest, *Nat. Photon.*, 2010, **4**, 371–375.
- 27 J.-W. Kang, B. Song, W. Liu, S.-J. Park, R. Agarwal and C.-H. Cho, *Sci. Adv.*, 2019, **5**, eaau9338.
- 28 E. Orgiu, J. George, J. A. Hutchison, E. Devaux, J. F. Dayen, B. Doudin, F. Stellacci, C. Genet, J. Schachenmayer, C. Genes, G. Pupillo, P. Samorì and T. W. Ebbesen, *Nat. Mater.*, 2015, **14**, 1123–1129.
- 29 X. Zhong, T. Chervy, L. Zhang, A. Thomas, J. George, C. Genet, J. A. Hutchison and T. W. Ebbesen, *Angew. Chem. Int. Ed.*, 2017, **56**, 9034–9038.
- 30 K. Georgiou, R. Jayaprakash, A. Othonos and D. G. Lidzey, *Angew. Chem. Int. Ed.*, 2021, **60**, 16661–16667.
- 31 M. Wang, M. Hertzog and K. Börjesson, *Nat. Commun.*, 2021, **12**, 1874.
- 32 D. M. Coles, N. Somaschi, P. Michetti, C. Clark, P. G. Lagoudakis, P. G. Savvidis and D. G. Lidzey, *Nat. Mater.*, 2014, **13**, 712–719.
- 33 M. Dusel, S. Betzold, O. A. Egorov, S. Klembt, J. Ohmer, U. Fischer, S. Höfling and C. Schneider, *Nat. Commun.*, 2020, **11**, 2863.
- 34 A. V. Zasedatelev, A. V. Baranikov, D. Urbonas, F. Scafirimuto, U. Scherf, T. Stöferle, R. F. Mahrt and P. G. Lagoudakis, *Nat. Photon.*, 2019, **13**, 378–383.
- 35 A. Kavokin, T. C. H. Liew, C. Schneider, P. G. Lagoudakis, S. Klembt and S. Höfling, *Nat. Rev. Phys.*, 2022, **4**, 435–451.
- 36 F. Pavošević, R. L. Smith and A. Rubio, *Catalysis in Click Chemistry Reactions Controlled by Cavity Quantum Vacuum Fluctuations: The Case of endo/exo Diels-Alder Reaction*, 2022, <http://arxiv.org/abs/2208.06925>, arXiv:2208.06925 [physics].

- 37 F. Pavošević, S. Hammes-Schiffer, A. Rubio and J. Flick, *J. Am. Chem. Soc.*, 2022.
- 38 C. Schäfer, J. Flick, E. Ronca, P. Narang and A. Rubio, *Nat. Commun.*, 2022, **13**, 7817.
- 39 C. Schäfer, *J. Phys. Chem. Lett.*, 2022, **13**, 6905–6911.
- 40 R. J. Cave and M. D. Newton, *J. Chem. Phys.*, 1997, **106**, 9213–9226.
- 41 L. A. Martínez-Martínez, R. F. Ribeiro, J. C. González-Angulo and J. Yuen-Zhou, *ACS Photonics*, 2017, **5**, 167–176.
- 42 P.-Y. Yang and J. Cao, *J. Phys. Chem. Lett.*, 2021, **12**, 9531–9538.
- 43 C. Climent and J. Feist, *Phys. Chem. Chem. Phys.*, 2020, **22**, 23545–23552.
- 44 D. S. Wang, T. Neuman, S. F. Yelin and J. Flick, *J. Phys. Chem. Lett.*, 2022, **13**, 3317–3324.
- 45 M. V. Imperatore, J. B. Asbury and N. C. Giebink, *J. Chem. Phys.*, 2021, **154**, 191103.
- 46 J. Galego, F. J. Garcia-Vidal and J. Feist, *Nat. Commun.*, 2016, **7**, 13841.
- 47 J. A. Campos-Gonzalez-Angulo, R. F. Ribeiro and J. Yuen-Zhou, *Nat. Commun.*, 2019, **10**, 4685.
- 48 J. P. Philbin, Y. Wang, P. Narang and W. Dou, *J. Phys. Chem. C*, 2022, **126**, 14908–14913.
- 49 S. Efrima and M. Bixon, *Chem. Phys. Lett.*, 1974, **25**, 34–37.
- 50 N. T. Phuc, P. Q. Trung and A. Ishizaki, *Sci. Rep.*, 2020, **10**, 7318.
- 51 E. Davidsson and M. Kowalewski, *J. Chem. Phys.*, 2020, **153**, 234304.
- 52 J. Galego, F. J. Garcia-Vidal and J. Feist, *Phys. Rev. Lett.*, 2017, **119**, 136001.
- 53 L. Mauro, K. Caicedo, G. Jonusauskas and R. Avriller, *Phys. Rev. B*, 2021, **103**, 165412.
- 54 I. Vurgaftman, B. S. Simpkins, A. D. Dunkelberger and J. C. Owrutsky, *J. Phys. Chem. Lett.*, 2020, **11**, 3557–3562.
- 55 H. Hiura and A. Shalabney, *ChemRxiv*, 2021.
- 56 H. Hiura, A. Shalabney and J. George, *ChemRxiv*, 2019.
- 57 D. Craig and T. Thirunamachandran, *Molecular Quantum Electrodynamics: An Introduction to Radiation-molecule Interactions*, Dover Publications, 1998.
- 58 A. Archambault, F. m. c. Marquier, J.-J. Greffet and C. Arnold, *Phys. Rev. B*, 2010, **82**, 035411.
- 59 D. L. Andrews, D. P. Craig and T. Thirunamachandran, *Int. Rev. Phys. Chem.*, 1989, **8**, 339–383.
- 60 A. W. Snyder and J. Love, *Optical Waveguide Theory*, Springer, 1st edn, 1983.
- 61 R. Chikkaraddy, B. de Nijs, F. Benz, S. J. Barrow, O. A. Scherman, E. Rosta, A. Demetriadou, P. Fox, O. Hess and J. J. Baumberg, *Nature*, 2016, **535**, 127–30.
- 62 F. Benz, M. K. Schmidt, A. Dreismann, R. Chikkaraddy, Y. Zhang, A. Demetriadou, C. Carnegie, H. Ohadi, B. de Nijs, R. Esteban, J. Aizpurua and J. J. Baumberg, *Science*, 2016, **354**, 726–729.
- 63 T. Neuman, R. Esteban, D. Casanova, F. J. García-Vidal and J. Aizpurua, *Nano Lett.*, 2018, **18**, 2358–2364.
- 64 D. S. Wang, T. Neuman and P. Narang, *J. Phys. Chem. C*, 2021, **125**, 6222–6228.
- 65 K. L. Kelly, E. Coronado, L. L. Zhao and G. C. Schatz, *J. Phys. Chem. B*, 2003, **107**, 668–677.
- 66 J. A. Schuller, E. S. Barnard, W. Cai, Y. C. Jun, J. S. White and M. L. Brongersma, *Nat. Mater.*, 2010, **9**, 193–204.
- 67 Jiang, K. Bosnick, M. Maillard and L. Brus, *J. Phys. Chem. B*, 2003, **107**, 9964–9972.
- 68 L. Brus, *Acc. Chem. Res.*, 2008, **41**, 1742–1749.
- 69 S. Nie and S. R. Emory, *Science*, 1997, **275**, 1102–1106.
- 70 C. Zhan, X.-J. Chen, J. Yi, J.-F. Li, D.-Y. Wu and Z.-Q. Tian, *Nat. Rev. Chem.*, 2018, **2**, 216–230.
- 71 H. A. Atwater and A. Polman, *Nat. Mater.*, 2010, **9**, 205–213.
- 72 Y. Zhang, C. Yam and G. C. Schatz, *J. Phys. Chem. Lett.*, 2016, **7**, 1852–1858.
- 73 R.-M. Ma, R. F. Oulton, V. J. Sorger and X. Zhang, *Laser Photonics Rev.*, 2013, **7**, 1–21.
- 74 P. Berini and I. De Leon, *Nat. Photon.*, 2012, **6**, 16–24.
- 75 J. Guan, R. Li, X. G. Juarez, A. D. Sample, Y. Wang, G. C. Schatz and T. W. Odom, *Adv. Mater.*, 2021, **n/a**, 2103262.
- 76 J. Guan, J.-E. Park, S. Deng, M. J. H. Tan, J. Hu and T. W. Odom, *Chem. Rev.*, 2022, **122**, 15177–15203.
- 77 S. K. Cushing and N. Wu, *J. Phys. Chem. Lett.*, 2016, **7**, 666–675.
- 78 J. Li, S. K. Cushing, F. Meng, T. R. Senty, A. D. Bristow and N. Wu, *Nat. Photon.*, 2015, **9**, 601–607.
- 79 Y. Zhang, *J. Phys. Chem. A*, 2021, **125**, 9201–9208.
- 80 C. Clavero, *Nat. Photon.*, 2014, **8**, 95–103.
- 81 M. Bernardi, J. Mustafa, J. B. Neaton and S. G. Louie, *Nat. Commun.*, 2015, **6**, 7044.
- 82 A. M. Brown, R. Sundararaman, P. Narang, I. William A. Goddard and H. A. Atwater, *ACS Nano*, 2016, **10**, 957–966.
- 83 A. Manjavacas, J. G. Liu, V. Kulkarni and P. Nordlander, *ACS Nano*, 2014, **8**, 7630–7638.
- 84 L. Yuan, J. Zhou, M. Zhang, X. Wen, J. M. P. Martirez, H. Robatjazi, L. Zhou, E. A. Carter, P. Nordlander and N. J. Halas, *ACS Nano*, 2022, **16**, 17365–17375.
- 85 L. Zhou, J. M. P. Martirez, J. Finzel, C. Zhang, D. F. Swearer, S. Tian, H. Robatjazi, M. Lou, L. Dong, L. Henderson, P. Christopher, E. A. Carter, P. Nordlander and N. J. Halas, *Nat. Energy*, 2020, **5**, 61–70.
- 86 L. Zhou, D. F. Swearer, C. Zhang, H. Robatjazi, H. Zhao, L. Henderson, L. Dong, P. Christopher, E. A. Carter, P. Nordlander and N. J. Halas, *Science*, 2018, **362**, 69–72.
- 87 X. Wu, T. v. d. Heide, T. Frauenheim, S. Tretiak, C. Yam and Y. Zhang, *arXiv Preprint*, 2023, **arXiv:2211.12625**, 0.
- 88 Y. Zhang, T. Nelson, S. Tretiak, H. Guo and G. C. Schatz, *Acs Nano*, 2018, **12**, 8415–8422.
- 89 Q. Wu, L. Zhou, G. C. Schatz, Y. Zhang and H. Guo, *J. Am. Chem. Soc.*, 2020, **142**, 13090–13101.
- 90 E. Kazuma, J. Jung, H. Ueba, M. Trenary and Y. Kim, *Science*, 2018, **360**, 521–526.

- 91 Y. Zhang, S. He, W. Guo, Y. Hu, J. Huang, J. R. Mulcahy and W. D. Wei, *Chem. Rev.*, 2018, **118**, 2927–2954.
- 92 T. E. Tesema, B. Kafle and T. G. Habteyes, *J. Phys. Chem. C*, 2019, **123**, 8469–8483.
- 93 J. R. Adleman, D. A. Boyd, D. G. Goodwin and D. Psaltis, *Nano Lett.*, 2009, **9**, 4417–4423.
- 94 C. Boerigter, R. Campana, M. Morabito and S. Linic, *Nat. Commun.*, 2016, **7**, 10545–10545.
- 95 Y. Zhang, T. Nelson and S. Tretiak, *Computational Photocatalysis: Modeling of Photophysics and Photochemistry at Interfaces*, ACS Publications, 2019, pp. 239–256.
- 96 D. Marinica, A. Kazansky, P. Nordlander, J. Aizpurua and A. G. Borisov, *Nano Lett.*, 2012, **12**, 1333–1339.
- 97 A. Varas, P. García-González, J. Feist, F. García-Vidal and A. Rubio, *Nanophotonics*, 2016, **5**, 409–426.
- 98 T. Perera, S. D. Gunapala, M. I. Stockman and M. Premaratne, *J. Phys. Chem. C*, 2020, **124**, 27694–27708.
- 99 T. P. Rossi, M. Kuisma, M. J. Puska, R. M. Nieminen and P. Erhart, *J. Chem. Theory Comput.*, 2017, **13**, 4779–4790.
- 100 N. Asadi-Aghbolaghi, R. Rüger, Z. Jamshidi and L. Visscher, *J. Phys. Chem. C*, 2020, **124**, 7946–7955.
- 101 P. Pandeya and C. M. Aikens, *J. Phys. Chem. A*, 2021, **125**, 4847–4860.
- 102 F. Alkan and C. M. Aikens, *J. Phys. Chem. C*, 2018, **122**, 23639–23650.
- 103 F. Alkan and C. M. Aikens, *J. Phys. Chem. C*, 2021, **125**, 12198–12206.
- 104 F. Della Sala, *J. Chem. Phys.*, 2022, **157**, 104101.
- 105 S. D'Agostino, R. Rinaldi, G. Cuniberti and F. Della Sala, *J. Phys. Chem. C*, 2018, **122**, 19756–19766.
- 106 J. W. You and N. C. Panoiu, *IEEE J. Multiscale and Multiphys. Comput. Techn.*, 2019, **4**, 111–118.
- 107 P. Hohenberg and W. Kohn, *Phys. Rev.*, 1964, **136**, B864–B871.
- 108 R. G. Parr and W. Yang, *Annu. Rev. Phys. Chem.*, 1995, **46**, 701–728.
- 109 E. Runge and E. K. U. Gross, *Phys. Rev. Lett.*, 1984, **52**, 997–1000.
- 110 M. E. Casida and M. Huix-Rotllant, *Annu. Rev. Phys. Chem.*, 2012, **63**, 287–323.
- 111 M. L. Tiago and J. R. Chelikowsky, *Phys. Rev. B*, 2006, **73**, 205334.
- 112 L. Román Castellanos, O. Hess and J. Lischner, *Commun. Phys.*, 2019, **2**, 47.
- 113 X. Wu, B. Liu, T. Frauenheim, S. Tretiak, C. Yam and Y. Zhang, *J. Chem. Phys.*, 2022, **157**, 214201.
- 114 J. G. Liu, H. Zhang, S. Link and P. Nordlander, *ACS Photonics*, 2018, **5**, 2584–2595.
- 115 H. Breuer, F. Petruccione and S. Petruccione, *The Theory of Open Quantum Systems*, Oxford University Press, 2002.
- 116 H. Haug and A. Jauho, *Quantum Kinetics in Transport and Optics of Semiconductors*, Springer Berlin Heidelberg, 1998.
- 117 T. E. Tesema, B. Kafle, M. G. Tadesse and T. G. Habteyes, *J. Phys. Chem. C*, 2017, **121**, 7421–7428.
- 118 C. Boerigter, U. Aslam and S. Linic, *ACS Nano*, 2016, **10**, 6108–6115.
- 119 R. Long and O. V. Prezhdo, *J. Am. Chem. Soc.*, 2014, **136**, 4343–4354.
- 120 R. Long, N. J. English and O. V. Prezhdo, *J. Am. Chem. Soc.*, 2012, **134**, 14238–14248.
- 121 T. Bora, D. Zoepfl and J. Dutta, *Sci. Rep.*, 2016, **6**, 26913.
- 122 Y. Sivan, I. W. Un and Y. Dubi, *Faraday Discuss.*, 2019, **214**, 215–233.
- 123 A. M. Brown, R. Sundararaman, P. Narang, A. M. Schwartzberg, W. A. Goddard and H. A. Atwater, *Phys. Rev. Lett.*, 2017, **118**, 087401.
- 124 T. R. Nelson, A. J. White, J. A. Bjorgaard, A. E. Sifain, Y. Zhang, B. Nebgen, S. Fernandez-Alberti, D. Mozyrsky, A. E. Roitberg and S. Tretiak, *Chem. Rev.*, 2020, **120**, 2215–2287.
- 125 J. C. Tully, *J. Chem. Phys.*, 1990, **93**, 1061–1071.
- 126 J. C. Tully, *J. Chem. Phys.*, 2012, **137**, 22A301.
- 127 B. F. Curchod, U. Rothlisberger and I. Tavernelli, *Chem. Phys. Chem.*, 2013, **14**, 1314–1340.
- 128 W. Malone, B. Nebgen, A. White, Y. Zhang, H. Song, J. A. Bjorgaard, A. E. Sifain, B. Rodriguez-Hernandez, V. M. Freixas, S. Fernandez-Alberti et al., *J. Chem. Theory Comput.*, 2020, **16**, 5771–5783.
- 129 E. Fabiano, T. W. Keal and W. Thiel, *Chem. Phys.*, 2008, **349**, 334–347.
- 130 D. V. Makhov, C. Symonds, S. Fernandez-Alberti and D. V. Shalashilin, *Chem. Phys.*, 2017, **493**, 200–218.
- 131 D. V. Makhov, W. J. Glover, T. J. Martinez and D. V. Shalashilin, *J. Chem. Phys.*, 2014, **141**, 054110.
- 132 D. V. Makhov, K. Saita, T. J. Martinez and D. V. Shalashilin, *Phys. Chem. Chem. Phys.*, 2015, **17**, 3316–3325.
- 133 D. V. Makhov, T. J. Martinez and D. V. Shalashilin, *Faraday Discuss.*, 2016, **194**, 81–94.
- 134 H. Song, V. M. Freixas, S. Fernandez-Alberti, A. J. White, Y. Zhang, S. Mukamel, N. Govind and S. Tretiak, *J. Chem. Theory Comput.*, 2021, **17**, 3629–3643.
- 135 M. Ben-Nun, J. Quenneville and T. J. Martínez, *J. Phys. Chem. A*, 2000, **104**, 5161–5175.
- 136 L. V. Besteiro, X.-T. Kong, Z. Wang, G. Hartland and A. O. Govorov, *ACS Photonics*, 2017, **4**, 2759–2781.
- 137 L. Chang, L. V. Besteiro, J. Sun, E. Y. Santiago, S. K. Gray, Z. Wang and A. O. Govorov, *ACS Energy Lett.*, 2019, **4**, 2552–2568.
- 138 L. Yan, F. Wang and S. Meng, *ACS Nano*, 2016, **10**, 5452–5458.
- 139 A. V. Akimov, A. J. Neukirch and O. V. Prezhdo, *Chem. Rev.*, 2013, **113**, 4496–4565.
- 140 F. Trani, G. Scalmani, G. Zheng, I. Carnimeo, M. J. Frisch and V. Barone, *J. Chem. Theory Comput.*, 2011, **7**, 3304–3313.
- 141 T. A. Niehaus, S. Suhai, F. Della Sala, P. Lugli, M. Elstner, G. Seifert and T. Frauenheim, *Phys. Rev. B*, 2001, **63**, 085108.

- 142 H. Song, S. A. Fischer, Y. Zhang, C. J. Cramer, S. Mukamel, N. Govind and S. Tretiak, *J. Chem. Theory Comput.*, 2020, **16**, 6418–6427.
- 143 X. Wu, S. Wen, H. Song, T. Frauenheim, S. Tretiak, C. Yam and Y. Zhang, *J. Chem. Phys.*, 2022, **157**, 084114.
- 144 B. Hourahine, B. Aradi, V. Blum, F. Bonafé, A. Buccheri, C. Camacho, C. Cevallos, M. Deshaye, T. Dumitrica, A. Dominguez *et al.*, *J. Chem. Phys.*, 2020, **152**, 124101.
- 145 O. A. Douglas-Gallardo, M. Berdakin, T. Frauenheim and C. G. Sánchez, *Nanoscale*, 2019, **11**, 8604–8615.
- 146 M. Berdakin, O. A. Douglas-Gallardo and C. G. Sanchez, *J. Phys. Chem. C*, 2019, **124**, 1631–1639.
- 147 J. A. Hutchison, T. Schwartz, C. Genet, E. Devaux and T. W. Ebbesen, *Angew. Chem. Int. Ed.*, 2012, **51**, 1592–1596.
- 148 J. George, T. Chervy, A. Shalabney, E. Devaux, H. Hiura, C. Genet and T. W. Ebbesen, *Phys. Rev. Lett.*, 2016, **117**, 153601.
- 149 R. M. A. Vergauwe, A. Thomas, K. Nagarajan, A. Shalabney, J. George, T. Chervy, M. Seidel, E. Devaux, V. Torbeev and T. W. Ebbesen, *Angew. Chem. Int. Ed.*, 2019, **58**, 15324–15328.
- 150 J. Lather, P. Bhatt, A. Thomas, T. W. Ebbesen and J. George, *Angew. Chem. Int. Ed.*, 2019, **58**, 10635–10638.
- 151 A. Sau, K. Nagarajan, B. Patrahanu, L. Lethuillier-Karl, R. M. A. Vergauwe, A. Thomas, J. Moran, C. Genet and T. W. Ebbesen, *Angew. Chem. Int. Ed.*, 2021, **60**, 5712–5717.
- 152 K. Hirai, H. Ishikawa, T. Chervy, J. A. Hutchison and H. Uji-i, *Chem. Sci.*, 2021, **12**, 11986–11994.
- 153 W. M. Takele, F. Wackenhut, Q. Liu, L. Piatkowski, J. Waluk and A. J. Meixner, *J. Phys. Chem. C*, 2021, **125**, 14932–14939.
- 154 A. Thomas, L. Lethuillier-Karl, K. Nagarajan, R. M. A. Vergauwe, J. George, T. Chervy, A. Shalabney, E. Devaux, C. Genet, J. Moran and T. W. Ebbesen, *Science*, 2019, **363**, 615–619.
- 155 J. Lather, A. N. K. Thabassum, J. Singh and J. George, *Chem. Sci.*, 2022, **13**, 195–202.
- 156 S. Tretiak and S. Mukamel, *Chem. Rev.*, 2002, **102**, 3171–3212.
- 157 J. Yang, Q. Ou, Z. Pei, H. Wang, B. Weng, Z. Shuai, K. Mullen and Y. Shao, *J. Chem. Phys.*, 2021, **155**, 064107.
- 158 Y. Zhang, T. Nelson and S. Tretiak, *J. Chem. Phys.*, 2019, **151**, 154109.
- 159 R. H. Tichauer, J. Feist and G. Groenhof, *J. Chem. Phys.*, 2021, **154**, 104112.
- 160 G. Groenhof and J. J. Toppari, *J. Phys. Chem. Lett.*, 2018, **9**, 4848–4851.
- 161 H. L. Luk, J. Feist, J. J. Toppari and G. Groenhof, *J. Chem. Theory Comput.*, 2017, **13**, 4324–4335.
- 162 G. Groenhof, C. Climent, J. Feist, D. Morozov and J. J. Toppari, *J. Phys. Chem. Lett.*, 2019, **10**, 5476–5483.
- 163 R. H. Tichauer, D. Morozov, I. Sokolovskii, J. J. Toppari and G. Groenhof, *J. Phys. Chem. Lett.*, 2022, **13**, 6259–6267.
- 164 A. Mandal, M. Taylor, B. Weight, E. Koessler, X. Li and P. Huo, *Theoretical Advances in Polariton Chemistry and Molecular Cavity Quantum Electrodynamics*, 2022, <https://chemrxiv.org/engage/chemrxiv/article-details/63903eee92f084c9612a9086>.
- 165 V. Rokaj, D. M. Welakuh, M. Ruggenthaler and A. Rubio, *J. Phys. B: At. Mol. Opt. Phys.*, 2018, **51**, 034005.
- 166 A. Mandal, T. D. Krauss and P. Huo, *J. Phys. Chem. B*, 2020, **124**, 6321–6340.
- 167 B. Weight, T. Krauss and P. Huo, *Investigating Molecular Exciton-Polaritons using Many-body Electronic Structure Theory with Cavity Quantum Electrodynamics*, 2023, <https://chemrxiv.org/engage/chemrxiv/article-details/63ba35271699ca2dd7e74772>.
- 168 J. Flick and P. Narang, *J. Chem. Phys.*, 2020, **153**, 094116.
- 169 D. S. Wang, T. Neuman, J. Flick and P. Narang, *J. Chem. Phys.*, 2021, **154**, 104109.
- 170 J. McTague and J. J. Foley, *J. Chem. Phys.*, 2022, **156**, 154103.
- 171 N. Vu, G. M. McLeod, K. Hanson and A. E. DePrince, *J. Phys. Chem. A*, 2022, **126**, 9303–9312.
- 172 T. S. Haugland, E. Ronca, E. F. Kjønstad, A. Rubio and H. Koch, *Phys. Rev. X*, 2020, **10**, 041043.
- 173 T. S. Haugland, C. Schäfer, E. Ronca, A. Rubio and H. Koch, *J. Chem. Phys.*, 2021, **154**, 094113.
- 174 A. E. DePrince, *J. Chem. Phys.*, 2022, **154**, 094112.
- 175 M. D. Liebenthal, N. Vu and A. E. DePrince, *J. Chem. Phys.*, 2022, **156**, 054105.
- 176 J. Flick, H. Appel, M. Ruggenthaler and A. Rubio, *J. Chem. Theory Comput.*, 2017, **13**, 1616–1625.
- 177 P. Vindel-Zandbergen, L. M. Ibele, J.-K. Ha, S. K. Min, B. F. E. Curchod and N. T. Maitra, *J. Chem. Theory Comput.*, 2021.
- 178 I. S. Lee, J.-K. Ha, D. Han, T. I. Kim, S. W. Moon and S. K. Min, *J. Comput. Chem.*, 2021, **42**, 1755–1766.
- 179 A. Abedi, N. T. Maitra and E. K. U. Gross, *Phys. Rev. Lett.*, 2010, **105**, 123002.
- 180 S. K. Min, F. Agostini, I. Tavernelli and E. K. U. Gross, *J. Phys. Chem. Lett.*, 2017, **8**, 3048–3055.
- 181 F. Agostini, S. K. Min, A. Abedi and E. K. U. Gross, *J. Chem. Theory Comput.*, 2016, **12**, 2127–2143.
- 182 S. K. Min, F. Agostini and E. K. U. Gross, *Phys. Rev. Lett.*, 2015, **115**, 073001.
- 183 J.-K. Ha and S. K. Min, *J. Chem. Phys.*, 2022, **156**, 174109.
- 184 P. Vindel-Zandbergen, S. Matsika and N. T. Maitra, *J. Phys. Chem. Lett.*, 2022, **13**, 1785–1790.
- 185 N. M. Hoffmann, H. Appel, A. Rubio and N. T. Maitra, *Eur. Phys. J. B*, 2018, **91**, 180.
- 186 T. G. Philbin, *Amer. J. Phys.*, 2014, **82**, 742–748.
- 187 R. R. Riso, T. S. Haugland, E. Ronca and H. Koch, *Nat. Commun.*, 2022, **13**, 1368.
- 188 A. Mandal and P. Huo, *J. Phys. Chem. Lett.*, 2019, **10**, 5519–5529.
- 189 R. R. Riso, T. S. Haugland, E. Ronca and H. Koch, *J. Chem. Phys.*, 2022, **156**, 234103.
- 190 C. Pellegrini, J. Flick, I. V. Tokatly, H. Appel and A. Rubio,

- Phys. Rev. Lett.*, 2015, **115**, 093001.
- 191 M. Ruggenthaler, J. Flick, C. Pellegrini, H. Appel, I. V. Tokatly and A. Rubio, *Phys. Rev. A*, 2014, **90**, 012508.
- 192 M. Ruggenthaler, N. Tancogne-Dejean, J. Flick, H. Appel and A. Rubio, *Nat. Rev. Chem.*, 2018, **2**, 1–16.
- 193 J. Flick, M. Ruggenthaler, H. Appel and A. Rubio, *Proc. Nat. Acad. Sci.*, 2015, **112**, 15285–15290.
- 194 T. E. Li, Z. Tao and S. Hammes-Schiffer, *J. Chem. Theory Comput.*, 2022, **18**, 2774–2784.
- 195 T. E. Li and S. Hammes-Schiffer, *Electronic Born-Oppenheimer Approximation in Nuclear-Electronic Orbital Dynamics*, 2023, <http://arxiv.org/abs/2301.04076>, arXiv:2301.04076 [physics].
- 196 U. Mordovina, C. Bungey, H. Appel, P. J. Knowles, A. Rubio and F. R. Manby, *Phys. Rev. Research*, 2020, **2**, 023262.
- 197 J. Fregoni, T. S. Haugland, S. Pipolo, T. Giovannini, H. Koch and S. Corni, *Nano Lett.*, 2021, **21**, 6664–6670.
- 198 T. G. Philbin, *American Journal of Physics*, 2014, **82**, 742–748.
- 199 A. F. White, Y. Gao, A. J. Minnich and G. K.-L. Chan, *J. Chem. Phys.*, 2020, **153**, 224112.
- 200 T. E. Li, B. Cui, J. E. Subotnik and A. Nitzan, *Annu. Rev. Phys. Chem.*, 2021, **73**, 43–71.
- 201 A. Taflove and S. C. Hagness, *Computational Electrodynamics: The Finite-Difference Time-Domain Method*, Artech House, 3rd edn, 2005.
- 202 C. Schäfer and G. Johansson, *Phys. Rev. Lett.*, 2022, **128**, 156402.
- 203 C. Schäfer, F. Buchholz, M. Penz, M. Ruggenthaler and A. Rubio, *Proc. Nat. Acad. Sci.*, 2021, **118**, e2110464118.
- 204 A. F. Peterson, S. L. Ray and R. Mittra, *IEEE Transactions on Antennas and Propagation*, 1998, **46**, 276–294.
- 205 M. S. Tame, K. R. McEnery, S. K. Ozdemir, J. Lee, S. A. Maier and M. S. Kim, *Nat. Phys.*, 2013, **9**, 329–340.
- 206 A. Gaita-Ariño, F. Luis, S. Hill and E. Coronado, *Nature Chemistry*, 2019, **11**, 301–309.
- 207 D. Sanvitto and S. Kéna-Cohen, *Nature Materials*, 2016, **15**, 1061–1073.
- 208 X. Zheng, F. Wang, C. Y. Yam, Y. Mo and G. Chen, *Phys. Rev. B*, 2007, **75**, 195127.
- 209 Y. Zhang, S. Chen and G. Chen, *Phys. Rev. B*, 2013, **87**, 085110.
- 210 Y. Zhang, C. Y. Yam and G. Chen, *J. Chem. Phys.*, 2013, **138**, 164121.
- 211 L. Chen, Y. Zhang, G. Chen and I. Franco, *Nat. Commun.*, 2018, **9**, 1–12.
- 212 T. Boolakee, C. Heide, A. Garzón-Ramírez, H. B. Weber, I. Franco and P. Hommelhoff, *Nature*, 2022, **605**, 251–255.