Applying the resonant-state expansion to realistic materials with frequency dispersion

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The dispersive resonant-state expansion, developed for an accurate calculation of the resonant states in open optical systems with frequency dispersion, is applied here to realistic materials, such as metallic nanoparticles and semiconductor microspheres. The material permittivity is determined by fitting the measured indices of refraction and absorption with a generalized Drude-Lorentz model containing a number of poles in the complex frequency plane. Each Drude or Lorentz pole generates an infinite series of resonant states. Furthermore, for small nanoparticles, each of these poles produces a distinct surface plasmon polariton mode. The evolution of these multiple surface modes with increasing radius traces the transition from the electrostatic limit to significant retardation and radiation. Treating the optical phonon range in a semiconductor microsphere, a reststrahlen band separating the resonant states is found. Considering a small energy range around the semiconductor band gap, the transition from absorption to gain is described by inverting the Lorentz pole weight, which results in the formation of lasing resonant states. Interestingly, the series of resonant states converging towards the absorption pole from the lower frequency side reshapes for a gain pole into a clockwise loop approaching the pole from the higher frequency side, being separated from a series spanning from low to high frequencies and containing the lasing modes.

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I. INTRODUCTION

Applying the concept of resonant states (RSs) [1,2] to optical systems is a mathematically rigorous way of treating their electromagnetic resonances. RSs are the eigenmodes of an optical system which can be found by solving Maxwell’s wave equation with outgoing wave boundary conditions. In open systems, the RS eigenfrequencies \( \omega_n \), numbered by the index \( n \), are generally complex, which physically reflects the fact that the energy leaks out of the system. For an isolated resonance in an optical spectrum of the system, the real part of the corresponding \( \omega_n \) determines the center frequency of the resonance and the imaginary part its half-width at half-maximum, with the quality factor of the resonance given by half of the ratio of real to imaginary part.

The complex eigenfrequencies of the RSs come along with a significantly different normalization of their vectorial wave functions as compared to Hermitian systems which have real eigenvalues. In particular, the standard normalization, given by the volume integral of the square modulus of the wave function, diverges for the RSs. The normalization of the RSs in electromagnetics has been found only recently in [3] where an exact analytical form of the RS normalization for an arbitrary dielectric system was provided. For systems with dispersion, a suited numerical normalization of the RSs was introduced in [4], where a perfectly matched layer (PML), frequently used in various electromagnetic numerical solvers, plays the role of an artificial absorber suppressing the problematic exponential divergence of the RS field at large distances from the system. This was followed by other numerical approaches to the normalization of RSs, eliminating its dependence on PMLs [5,6], for systems with or without frequency dispersion. The exact analytic normalization (without using PMLs) was developed for systems with frequency dispersion of the permittivity in [7] and later on generalized for dispersion of the permeability and chirality [8]. For more details on the development of the normalization, see [7,9,10].

One way to avoid the spatial divergence related to the complex eigenfrequencies of the RSs is to treat resonances in optical spectra in terms of some eigenstates introduced for a fixed real frequency \( \omega \). A number of approaches were suggested in this direction, ranging from constant flux states originally introduced by Kapur and Peierls [11] to eigenpermittivity modes proposed by Bergman [12]. The concept of constant flux states has been intensively exploited for random dielectric lasers [13]. Constant flux states present a complete set of functions suited for field expansion within the systems. They also possess an orthonormality similar to that valid for Hermitian systems. However, the major disadvantage of this approach is that the constant flux states are not solutions of Maxwell’s equations, which makes them somewhat abstract mathematical objects, useful only for treating RSs with frequencies close to the chosen real \( \omega \). A similar idea has been recently presented in [14] with another eigenvalue problem introduced to replace Maxwell’s equations.

A more rigorous approach of this kind is the one based on eigenpermittivity states [12] which are solutions of Maxwell’s equations for a fixed real frequency. The eigenpermittivities are complex, given by the geometry only, and are not matching the permittivity of the considered structure. A recent work [15] presented an accurate calculation of the full dyadic Green’s function of plasmonic nanoparticles in two dimensions using its expansion into eigenpermittivity states. This demonstrated that the approach is promising and suited for systems with any frequency dispersion, at a fixed real frequency, such as a light scattering problem at a given excitation frequency.
Using instead the RSs permits much faster calculations of optical spectra, since the dependence on the frequency is given analytically by a single set of states: Any observable has the form of a superposition of complex Lorentzian terms, each term due to an individual RS. Such a superposition is at the heart of the resonant-state expansion (RSE) [3] and is mathematically based on applying the Mittag-Leffler theorem [16] to the dyadic Green’s function, which was demonstrated in [3,17,18] for one-, two-, and three-dimensional dielectric systems. A phenomenological approach to the normalization in a plasmonic system, applied for a few important RSs calculated numerically using PMLs, was introduced in [4], where an observable—the Purcell factor of a plasmonic system—was calculated as an expansion into these RSs. A rigorous theory of the Purcell effect using an arbitrary number of RSs and their exact normalization has then been presented in [7]. The PML approach [4] has been refined later on in [19] by including in the Green’s function expansion more RSs as well as PML modes, which is required for completeness. This effectively reduces the physical open optical system to a somewhat different closed one including the PML. The price paid for this is that (i) the majority of the eigenstates of this system are strongly affected by the PML, in this way deviating from the RSs of the original system, and (ii) there emerge a large number of PML modes carrying no physical meaning. A different approach, called Riesz-projection method, has been recently suggested in [20] for an efficient treatment of plasmonic systems with only a few RSs close to the frequency range of interest. It introduces a finite closed contour in the complex frequency plane, and numerically evaluates the contour integral. This idea can be understood as a modification of the Mittag-Leffler theorem, by using the residue theorem applied for a finite contour. We note that [20] introduces a cut of the GF with a continuum of artificial modes on the cut, due to its formulation on the $\omega^2$ plane. Even though the contour of integration can typically be chosen such that it avoids the cut, this complication could be removed by using the $\omega$ plane instead.

An overview of some approaches in plasmonics based on RSs has been given in [21]. The review highlights the major challenge for any numerical mode solver dealing with dispersion, which is that the frequency dispersion makes Maxwell’s equations nonlinear in frequency. One way of solving this problem is the auxiliary-field approach [19], linearizing Maxwell’s equations by introducing auxiliary fields of polarization and magnetization. The number of auxiliary fields grows linearly with the number of poles in the generalized Drude-Lorentz model of the permittivity [22,23]. A numerical linearization of a polynomial eigenvalue problem of Maxwell’s equation with the Drude-Lorentz dispersion was considered in [21] as an alternative. The review does not mention that the problem of nonlinearity of Maxwell’s equation with dispersion had been already solved in the approach called the dispersive RSE, by introducing alternative representations of the GF [22]. These representations, based on sum rules for RSs imposed by the dispersion, linearize the eigenvalue problem of Maxwell’s equation on a fundamental level. Furthermore, the RSE is a rigorous approach which is not using PMLs to simulate open space.

The RSE is a rigorous perturbative method developed in [3] for calculation of the RSs of an arbitrary finite optical system (called perturbed system). The RSE converts the Maxwell wave equation into a linear matrix eigenvalue problem using the RSs of a simpler and usually analytically solvable system (called unperturbed system) as a basis for expansion. In this way, the RSE is not limited to small perturbations and finds numerically exactly all the eigenmodes of the perturbed system. The RSE has been generalized [22] to optical systems with frequency dispersion, such as metal or semiconductor nanoparticles. This dispersive RSE (dRSE) is suited for materials with dispersion of the permittivity described by a generalized Drude-Lorentz model [23]. In practice, one can find the parameters of this model—generalized conductivities and complex poles of the permittivity—by fitting the measured indices of refraction and absorption, as we have recently demonstrated for metals [23] and semiconductors [24].

The RSE can treat perturbations of arbitrary strength [3], and can be superior to existing computational methods in electrodynamics, such as finite difference in time domain (FDTD) and finite element method (FEM), in terms of accuracy and efficiency [18,25]. Furthermore, a significant advantage of the RSE method is that it calculates the full spectrum of RSs in a wide frequency range striving towards completeness, and it does not depend on further approximations, i.e., it is numerically exact. These properties are retained in the dRSE. Indeed, the only parameter of the RSE is the number of the RSs included in the basis, which can in principle be made arbitrarily large. While the formalism of the RSE is analytically advanced, its technical implementation is rather straightforward, as it reduces solving Maxwell’s wave equation to a diagonalization of a matrix containing a perturbation. For the latter, one needs a complete set of RSs of an unperturbed system, ideally chosen in such a way that the matrix elements of the perturbation can be efficiently computed.

In the present work we apply the dRSE to metal and semiconductor nanoparticles, using as a basis system a nondispersive dielectric sphere. This approach has been used in Ref. [22] for a dispersion described by the Drude model. Here we extend the method to systems with a dispersion of the permittivity described by up to four pairs of Lorentz poles. Including these poles in the dispersion leads to new insights into the properties of metal and semiconductor particles, such as a coexistence of multiple surface plasmon polariton modes, which is of significance for the field of plasmonics [26]. We also explore how the dRSE can describe the onset of lasing when material absorption changes to gain in semiconductor microstructures, important for the field of optical microcavities [27,28] and semiconductor lasers [29].

The paper is organized as follows. In Sec. II we outline the formalism of the dRSE, providing general expressions for the normalization of dispersive RSs and for a linear matrix eigenvalue problem determining the perturbed resonant states. The dRSE formalism is based on the condition that the dispersion of the unperturbed and perturbed systems has the same pole structure. In order to allow the unperturbed system to have no dispersion, or the perturbation to introduce additional poles in the dispersion, we present in Sec. III a variation of the dRSE called infinitesimal dispersive RSE (idRSE), which extends the basis of the unperturbed RSs to include a subset...
of RSs, called pole RSs, created by the additional poles of the permittivity in the limit of infinitesimal conductivities. We then apply in Sec. IV the dRSE to metal and semiconductor spherically symmetric systems. Using the idRSE, we find in Sec. IV A the RSs of a gold sphere described by the Drude-Lorentz model with three pairs of Lorentz poles. This is done using the RSs of a dielectric sphere as a basis extended to include the pole RSs. We concentrate our study on the surface plasmon polariton RSs and investigate their dependence on the sphere radius changing from 1 to 100 nm. In Sec. IV B we consider the optical-phonon range of the permittivity of GaAs, which can be accurately fitted by one pair of Lorentz poles, and find the RSs of a 50-μm-radius GaAs microsphere. We calculate the lasing modes of GaAs microspheres with 10 and 1 μm radius, having the permittivity close to the GaAs band edge fitted by four pairs of classical Lorentz poles. The optical gain is introduced by inverting the weight of the pole at the band edge from absorption to gain. Details of the permittivity models and the calculation of basis RSs and their matrix elements are given in Appendices A–C.

II. FORMALISM OF THE DISPERSIVE RSE

The basis for the dRSE is formed from the RSs of an unperturbed system which are the eigensolutions of the Maxwell wave equation,

\[ \nabla \times \nabla \times \mathbf{E}_n(\mathbf{r}) = \frac{\omega^2}{c^2} \hat{\mathbf{e}}(\mathbf{r}, \omega_0) \mathbf{E}_n(\mathbf{r}), \]

solved with outgoing wave boundary conditions (here \( c \) is the speed of light in vacuum). The permittivity tensor \( \hat{\mathbf{e}}(\mathbf{r}, \omega) \) of the unperturbed system is taken as an analytic function in the complex frequency plane, with a countable number of simple poles. According to the Mittag-Leffler theorem [16], the frequency dispersion of the permittivity tensor can be expressed as

\[ \hat{\mathbf{e}}(\mathbf{r}, \omega) = \hat{\mathbf{e}}_\infty(\mathbf{r}) + \sum_j \frac{i\delta j(\mathbf{r})}{\omega - \Omega j}, \]

where \( \hat{\mathbf{e}}_\infty(\mathbf{r}) \) is the high-frequency value of the permittivity and \( \Omega j \) are the resonance frequencies, at which the permittivity has simple poles, with the residues at the poles given by generalized conductivity tensors \( \delta j(\mathbf{r}) \). Equation (2) is known as the generalized Drude-Lorentz model [23]; more details about the model and its special cases (Drude and Lorentz models) are provided in Appendix A.

The dRSE solves the Maxwell wave equation

\[ \nabla \times \nabla \times \mathbf{E}(\mathbf{r}) = \frac{\omega^2}{c^2} \left[ \hat{\mathbf{e}}(\mathbf{r}, \omega) + \Delta \hat{\mathbf{e}}(\mathbf{r}, \omega) \right] \mathbf{E}(\mathbf{r}) \]

for the perturbed system (the system of interest) described by a modified permittivity tensor \( \hat{\mathbf{e}}(\mathbf{r}, \omega) + \Delta \hat{\mathbf{e}}(\mathbf{r}, \omega) \), in which the perturbation

\[ \Delta \hat{\mathbf{e}}(\mathbf{r}, \omega) = \Delta \hat{\mathbf{e}}_\infty(\mathbf{r}) + \sum_j \frac{i\Delta \delta j(\mathbf{r})}{\omega - \Omega j} \]

has the same pole structure as \( \hat{\mathbf{e}}(\mathbf{r}, \omega) \), i.e., its poles are at the same frequencies \( \Omega j \) and tensors \( \Delta \delta j(\mathbf{r}) \) and \( \Delta \hat{\mathbf{e}}(\mathbf{r}, \omega) \) are frequency independent. The electric field \( \mathbf{E}(\mathbf{r}) \) and the frequency \( \omega \) in Eq. (3) are, respectively, the wave function and the eigenfrequency of a RS of the perturbed system. The perturbed Maxwell’s wave equation (3) can be solved with the help of the Green’s function (GF) \( \hat{\mathbf{G}}_m(\mathbf{r}, \mathbf{r}') \) of the unperturbed Eq. (1) in the following way:

\[ \mathbf{E}(\mathbf{r}) = \frac{-\omega^2}{c^2} \int \hat{\mathbf{G}}_m(\mathbf{r}, \mathbf{r}') \Delta \hat{\mathbf{e}}(\mathbf{r}', \omega) \mathbf{E}(\mathbf{r}') d\mathbf{r}'. \]

The completeness of the basis of the unperturbed RSs inside the basis sphere allows us to convert this integral equation to a matrix eigenvalue problem. This can be done by expanding the perturbed RS field into the unperturbed ones,

\[ \mathbf{E}(\mathbf{r}) = \sum_n c_n \mathbf{E}_n(\mathbf{r}), \]

and by using the Mittag-Leffler theorem again, this time for the Green’s dyadic,

\[ \hat{\mathbf{G}}_m(\mathbf{r}, \mathbf{r}') = c^2 \sum_n \frac{\mathbf{E}_n(\mathbf{r}) \otimes \mathbf{E}_n(\mathbf{r}')}{\omega \omega - \omega_n}. \]

Here \( \otimes \) denotes the dyadic product of the vector fields, and the RS wave functions \( \mathbf{E}_n(\mathbf{r}) \) are suitably normalized for this expansion [7,9,18,22]. For nonstatic RSs, i.e., those having \( \omega_n \neq 0 \), the correct normalization linked to Eq. (7) is given by [7,8,22]

\[ 1 = 2 \int_V \mathbf{E}_n(\mathbf{r}) \cdot \frac{\partial (\omega^2 \hat{\mathbf{e}}(\mathbf{r}, \omega))}{\partial (\omega^2)} \bigg|_{\omega_n} d\mathbf{r} \]

\[ + c^2 \int_{S_V} \left( \frac{\partial \mathbf{E}_n}{\partial s} - \mathbf{E}_n \frac{\partial \mathbf{E}_n}{\partial s} \right) dS, \]

where \( \mathbf{E}_n = (\mathbf{r} \cdot \nabla) \mathbf{E}_n \), \( V \) is an arbitrary volume containing the system, the surface \( S_V \) is its boundary, and \( \partial / \partial s \) is the spatial derivative along the outer surface normal. Note that the electric field normalized according to Eq. (8) is a factor of \( \sqrt{2} \) smaller than the one used in the previous works [3,22]. This change has been introduced in order to symmetrize the normalization with respect to the electric and magnetic field, as discussed in Ref. [8].

The sum in Eq. (7) is taken over all RSs; their eigenfrequencies \( \omega_n \) play the role of simple poles of the GF. Owing to the frequency dispersion Eq. (2), the GF has alternative spectral representations linked to Eq. (7) via sum rules which the RSs of the dispersive system obey [8,22]. Using in Eq. (5) a suited combination of the different spectral representations of the GF and equating the coefficients at the basis functions \( \mathbf{E}_n(\mathbf{r}) \), this integral equation reduces to the following matrix eigenvalue problem, linear in frequency \( \omega \):

\[ \omega_n \sum_m (\delta_{nm} - U_{nm}) c_m = \omega \sum_m (\delta_{nm} + V_{nm}) c_m, \]

which presents the key equation of the dRSE. Here \( V_{nm} \) and \( U_{nm} \) are the matrix elements of the nondispersive and dispersive parts of the perturbation, given by

\[ V_{nm} = \int \mathbf{E}_n(\mathbf{r}) \cdot \Delta \hat{\mathbf{e}}_\infty(\mathbf{r}) \mathbf{E}_m(\mathbf{r}) d\mathbf{r}, \]
and

\[ U_{nm} = \int \mathbf{E}_n(r) \cdot [\Delta \hat{\varepsilon}(r, \omega_n) - \Delta \hat{\varepsilon}_\infty(r)] \mathbf{E}_m(r) \, dr \tag{11} \]

\[ = \frac{i}{\omega_n - \Omega_j} \int \mathbf{E}_n(r) \cdot \Delta \hat{\sigma}_j(r) \mathbf{E}_m(r) \, dr, \tag{12} \]

respectively. For a detailed derivation of the dRSE equation (9), see Refs. [8,22].

III. INFINITESIMAL-DISPERSIVE BASIS

The dRSE formally requires the permittivity of the basis system and the perturbed system to have the same poles. While this appears as a limitation of the dRSE, we can treat new poles of the perturbation \( \Delta \hat{\varepsilon}(r, \omega) \) by simply adding them to the basis system with negligible conductivities. This situation is illustrated in Sec. IV, where an unperturbed system without frequency dispersion, such as a dielectric sphere in vacuum, is used as a basis for the dRSE treating a system with dispersion, such as a metal or a semiconductor nanoparticle.

In the functional space of dispersive solutions, the basis of nondispersive RSs is incomplete and needs to be complemented (for completeness) with so-called pole RSs (pRSs). These pRSs originate from a group of RSs of a dispersive system with a permittivity pole at \( \omega = \Omega_j \) having an infinitesimal conductivity \( \hat{\sigma}_j \to 0 \). All the pRSs due to this pole then become degenerate, \( \omega_n \to \Omega_j \), while having different values of the effective refractive index, and hence different wave functions. These wave functions, however, have vanishing normalization constants \( A_n \to 0 \), so that without perturbing the pole to a finite conductivity, the pRSs do not contribute to the modes of the system, as both \( V_{nm} \) and \( U_{nm} \) are vanishing for \( n \) or \( m \) referring to pRSs.

To treat the pRSs of the \( j \)th pole of the permittivity, we introduce in the basis system an infinitesimal conductivity \( \Delta \hat{\sigma}_j(r) = \xi j_j(r) \) with \( \xi \to 0 \), \( \hat{\sigma}_j(r) \) is an arbitrary finite tensor, which can be chosen, without loss of generality, as \( \hat{\sigma}_j(r) = \mathbf{1} \) within the volume \( V \) of the system and zero outside it. In the limit \( \xi \to 0 \), the eigenfrequencies of the pRSs become

\[ \omega_n = \Omega_j + \xi q_n, \tag{13} \]

where \( q_n \) are finite eigenvalues determined by the geometry of the system and its material parameters. For example, for transverse-magnetic (TM) modes of a sphere in vacuum they are found solving a secular equation (B8), see Appendix B. Note that for brevity of notations, here and below we have omitted index \( j \) which would appear in all the eigenvalues and matrix elements due to the pRSs of pole \( j \).

According to Eqs. (2) and (13), the permittivity calculated at the pRS frequency \( \omega_n \), contributing both to Eqs. (8) and (11), is given by

\[ \hat{\varepsilon}(r, \omega_n) = \hat{\varepsilon}_\infty(r) + \sum_{j \neq j} \frac{i \hat{\sigma}_j(r)}{\Omega_j - \Omega_j'} + \xi j_j(r) q_n, \tag{14} \]

which is a finite quantity, determining the effective refractive index and hence the wavelength of the pRS. However, the frequency derivative of the permittivity, contributing to the RS normalization Eq. (8), is divergent,

\[ \frac{\partial \hat{\varepsilon}(r, \omega)}{\partial \omega} \bigg|_{\omega_n} = -\frac{1}{\Omega_j} \xi j_j(r), \tag{15} \]

which leads to a vanishing normalization coefficient of the pRS field, as follows from Eq. (8):

\[ \xi = -\frac{\Omega_j}{q_n^2} \int \mathbf{E}_n(r) \cdot \hat{\sigma}_j(r) \mathbf{E}_m(r) \, dr, \tag{16} \]

since \( \xi \to 0 \) but \( \Omega_j/q_n^2 \) is finite.

For any finite perturbation of the conductivity \( \Delta \hat{\sigma}_j \neq 0 \), the vanishing normalization constants of pole RSs are compensated in the perturbation matrix by divergencies due to the dispersion \( i \Delta \hat{\sigma}_j/(\omega_n - \Omega_j) \propto 1/\xi \), thus making \( U_{nm} \) finite. To take this limit properly, it is convenient to introduce the factors

\[ \alpha_n = \left\{ \begin{array}{ll} 1 & \text{for non-pRSs,} \\ \Omega_j/\Omega_j' & \text{for pRSs of } \Omega_j, \end{array} \right. \tag{17} \]

and to redefine the RS wave functions

\[ \mathbf{E}_n(r) = \alpha_n \tilde{\mathbf{E}}_n(r), \tag{18} \]

where all \( \tilde{\mathbf{E}}_n(r) \) have finite normalization. Expressing the perturbation matrices and eigenvectors as

\[ V_{nm} = \alpha_n \alpha_m Q_{nm}, \quad U_{nm} = \frac{\alpha_m}{\alpha_n} S_{nm}, \quad c_n = \frac{b_n}{\alpha_n}, \tag{19} \]

d the dRSE matrix equation (9) takes the form

\[ \alpha_n \sum_m (\delta_{nm} - S_{nm}) b_m = \omega \sum_m (\delta_{nm} + \alpha^2_n Q_{nm}) b_m, \tag{20} \]

in which the elements \( Q_{nm} \) and \( S_{nm} \) are finite. Explicitly, the matrix \( Q_{nm} \) is defined according to

\[ Q_{nm} = \int \tilde{\mathbf{E}}_n(r) \cdot \Delta \hat{\varepsilon}_\infty(r) \tilde{\mathbf{E}}_m(r) \, dr, \tag{21} \]

while the matrix \( S_{nm} \) is given by

\[ S_{nm} = \int \tilde{\mathbf{E}}_n(r) \cdot [\Delta \hat{\varepsilon}_n(r, \omega_n) - \Delta \hat{\varepsilon}_\infty(r)] \tilde{\mathbf{E}}_m(r) \, dr, \tag{22} \]

if \( n \) refers to a non-pRS, and by

\[ S_{nm} = \frac{i}{\Omega_j} \int \tilde{\mathbf{E}}_n(r) \cdot \Delta \hat{\sigma}_j(r) \tilde{\mathbf{E}}_m(r) \, dr, \tag{23} \]

if \( n \) refers to a pRS with \( \omega_n = \Omega_j \). Accordingly, the perturbed field is given by

\[ \mathbf{E}(r) = \sum_n b_n \tilde{\mathbf{E}}_n(r) \tag{24} \]

in terms of the modified basis RS wave functions \( \tilde{\mathbf{E}}_n(r) \) and the eigenvector components \( b_n \) determined by Eq. (20). Clearly, if \( \hat{\sigma}_j(r) = \Delta \hat{\sigma}_j(r) = 0 \), none of the corresponding pRSs is expected to contribute to the perturbed field. Indeed, in this case \( S_{nm} = 0 \) for any \( j \)th pRS \( n \), according to Eq. (23), while \( \alpha_n = 0 \) is leading to \( \alpha^2_n Q_{nm} = 0 \) in Eq. (20), even though \( V_{nm} \neq 0 \). Hence \( b_n = 0 \), showing that this pRS does not contribute to the perturbed RSs.
The method presented in this section is called infinitesimal-dispersive RSE (idRSE). The described procedure is suited for any number of new poles introduced in the perturbation. It has been already applied to systems with dispersion described by the Drude model [22]. In Sec. IV below we illustrate the idRSE on several examples, for the dispersion of the permittivity including both Drude and Lorentz poles, or Lorentz poles only.

IV. RESULTS

In this section we present a selection of applications of the dRSE. To reiterate, the requirements to apply the dRSE are: (i) the permittivity is described by the DL model, (ii) the perturbed system is enclosed within the basis sphere, and (iii) all basis RSs fulfilling the frequency cutoff condition are found. Even though the dRSE can treat arbitrary spatial dependencies of \( \hat{\sigma}_j(r) \) and \( \hat{\varepsilon}_\infty(r) \) within the basis sphere, in the present work we treat only homogeneous perturbations within the sphere, as we focus on the effect of changing the material dispersion. This also allows us to compare results of the dRSE with available exact solutions.

We consider TM polarization, and treat \( l = 1 \) modes of metallic particles, showing localized surface plasmons (SPs), where \( l \) is the orbital quantum number. For semiconductor microspheres instead, we use \( l \gg 1 \) in order to obtain sharp whispering-gallery (WG) modes. The wave functions, normalization coefficients, and the secular equation determining the RS eigenvalues of a uniform sphere in vacuum are provided in Appendix B, along with the matrix elements for uniform perturbation. Approximate recursive relations determining the start values for solving the secular equation are given in Appendix C. The basis for the dRSE is truncated in such a way that all unperturbed RSs satisfying the condition \( \{n_i(\omega_0)\omega_0\}_1 < \Omega_c \) are included in the basis, where \( n_i(\omega) = \sqrt{\varepsilon(\omega)} = n_i' + i n_i'' \) is the complex refractive index, and \( \Omega_c \) is the cut-off frequency determining the basis size \( N \). For each \( l \), a longitudinal static mode [18] with \( \omega_0 = 0 \) has been included in the basis; this is required for completeness of a nondispersive basis or a basis without Ohm’s law pole in the dispersion.

To apply the dRSE to spherical particles made of realistic materials, such as gold and GaAs considered below, we have generated the corresponding parameters of the Drude-Lorentz model using the earlier developed fit program [23]. This fit program is using the experimentally measured indices of refraction and absorption, \( n_i' \) and \( n_i'' \), respectively, in order to find the optimal values of the pole frequencies \( \Omega_j \) and their associated conductivities \( \sigma_j \). The value of the high-frequency permittivity \( \varepsilon_\infty \) can be used as an adjustable parameter which can be chosen to generate a stable system. The basis system corresponds to the slowest convergence as in this case the refractive-index contrast between the system and the environment (vacuum) is minimized for high-frequency modes. In the example provided in Sec. IV A we transform a dielectric (silica) sphere with \( \varepsilon_\infty = 2.1272 \) to a gold sphere with \( \varepsilon_\infty + \Delta \varepsilon_\infty = 0.5 \), while in Secs. IV B–IV D we use \( \varepsilon_\infty = 11.0 \) or 8.6013 for calculating the RSs of a GaAs microsphere, while keeping \( \Delta \varepsilon_\infty = 0 \). Table I of Appendix A shows all the fit parameters of gold and GaAs used in the dRSE calculations presented in this section below.

A. Infinitesimal-dispersive RSE: Plasmons

To construct a basis for the idRSE treating a gold nanosphere, we add to the set of the RSs of a nondispersive dielectric nanosphere with \( n_r = 1.4585 \) the pRSs of a Drude pole \( \Omega_0 = -i \gamma \) and of three pairs of Lorentz poles \( \Omega_j = (j = \pm 1, \pm 2, \pm 3) \) with infinitesimal conductivities. The poles are shown in Fig. 1(b), lined up with the corresponding fit of \( n_i' \) and \( n_i'' \) of gold [30], which is illustrated in Fig. 1(a).
The spectrum of the perturbed system—a gold sphere of radius $R = 10$ nm—shown in Fig. 1(b) reveals RSs to the high energy side of each pole in the dispersion. These are known as surface plasmon (SP) polariton resonances. In the electrostatic limit, they occur at $\varepsilon(\omega) = -2$ for $l = 1$, as given by Eq. (B9) of Appendix B, and are well known in the literature [31]. This equation provides one SP state for each pole in $\varepsilon$. Accordingly, using the Drude model, one only such RS is present [22]. Notably, the RSE calculates these poles beyond the electrostatic limit—the resulting evolution of the SPs versus radius is discussed below. The validity of the response predicted by the SPs at complex frequencies depends on the reliability of the DL model of $\varepsilon(\omega)$ at these frequencies, keeping in mind that the parameters of the model are fitted to describe the permittivity at real frequencies. At small imaginary parts of the SP frequencies, such as for SP0, the validity is expected to be better than at larger imaginary parts, such as for SP1 and SP2. Importantly, for correct calculation of the response of the system at real frequencies, such as the scattering and absorption spectra, the complete set of its RSs is required [32], which necessarily includes all the SP modes generated by the model of $\varepsilon(\omega)$ used. The absorption line shape is therefore not just a single resonance, but is more complex, as also observed in the experiment.

In addition to the SP mode around each pole of the permittivity, the full solution of the secular equation [see Eq. (B8) of Appendix B] contains a countable infinite number of modes in the vicinity of the Drude and each Lorentz pole, originating from the prRSs. The inset in Fig. 1(b) shows these RSs approaching the first Lorentz pole $\Omega_1$. We can also see from the figure that the idRSE matches the exact perturbed solution; however, in the presence of Lorentz poles, it is difficult to find all modes within a frequency range reliably from the secular equation (B8). The idRSE can find these solutions due to the completeness of RSs and is expected to work equally well for systems without known analytic solutions [18]. The relative error of the idRSE, given in Fig. 1(c) for the SPs, shows a $N^{-3}$ dependence on the number of states in the basis, as observed previously for spherically symmetric systems [3,22].

The evolution of the SP modes with nanosphere radius $R$ changing from 1 to 100 nm is given in Fig. 2. We find that for $R < 15$ nm, the SP modes keep close to the electrostatic limit, as expected for sizes much smaller than the light wavelength, $kR/c \ll 1$, noting that the SP wavelengths are in the 250 to 500 nm range. Increasing $R$, different SP modes change differently. Generally they shift to longer wavelength, as expected from modes determined by size rather than dispersion. SP0, related to the Drude pole and thus representing the classical SP of gold, is not changing much in the full range of the radii shown, which is due to the dominant role of the Drude part in the dispersion of gold and a rather small value of $\gamma$ for the Drude pole, see Table I. For small $R$, SP0 has an energy around 2.4 eV and a linewidth of about 0.3 eV, comparable to the features seen in the measured absorption [31]. Increasing $R$, SP0 moves along a small arc, first shifting to the red, then increasing the magnitude of the imaginary part of the wave number due to the increasing role of the radiative broadening, and then shifting back to the blue, possibly due to a repulsion from SP1, the next closest surface plasmon mode originating from the Lorentz pole at $\Omega_1$. This SP mode, SP1, has an energy of 2.9 eV in the static regime and a linewidth of 1.5 eV; this mostly accounts for the experimentally observed intraband absorption features. Note however that in calculating the absorption and scattering properties, the SPs will interfere [32], so that the line shape is expected to be non-Lorentzian, again in agreement with experiment. Calculating these properties using the dRSE will be the topic of a future work. With increasing $R$, SP1 is approximately keeping its linewidth, but is redshifting, passing SP0 at around 80 nm. This leads to a redshift and broadening of the observed SP resonance in absorption, in qualitative agreement with experiment [33]. SP2 shows a similar behavior as SP1, but with a twice broader linewidth, therefore describing a broad absorption in the interband absorption range of gold.

B. Infinitesimal-dispersive RSE: Optical phonons

The optical phonons in GaAs provide a sharp resonance in the permittivity over the energy range of 28–40 meV. This is well described by a single Lorentz pole with a purely imaginary positive conductivity, corresponding to a classical Lorentz oscillator having a purely real negative weight of the permittivity pole. The refractive and absorption index spectra and their fit are shown in Fig. 3(a). The fit parameters are given in Table I of Appendix A. We note that from about 33 to 36 meV, the absorption index supersedes the refractive index, which is the so-called reststrahlen region where a negative real part of the permittivity is causing an increased reflectivity of the material.

We use the idRSE with a basis system given by a dielectric microsphere with the same $\varepsilon_\infty$ as used in the above fit. We choose $l = 15$ and $R = 50 \mu$m in order for the Lorentz pole $\Omega_1$ to be close to WG mode frequencies. The RS frequencies of the basis system and of the perturbed system calculated via idRSE and exactly via Eq. (B8), are shown in Fig. 3(c). Due to the large $l$, the basis is clearly separated into WG, Fabry-Perot (FP), and leaky RSs. In the perturbed system we find the WG RSs strongly modified. There is a series of RSs approaching $\Omega_1$ from the lower frequency side, see also a zoom in Fig. 3(b). This series is countable infinite, limited
FIG. 3. (a) Measured refractive index \( n' \) and absorption index \( n'' \) of GaAs in the optical phonon range [34] (circles) and their fit with the Lorentz model Eq. (A1) with \( L = 1 \) (solid lines) as functions of the photon energy \( \hbar \omega \). (b) RSs for \( l = 15 \) and TM polarization of the basis sphere of radius \( R = 50 \mu \text{m} \) with \( n_r = 3.317 \) (open circles) and of the perturbed sphere with the GaAs optical phonon resonance perturbation showing the Lorentz pole \( \Omega_1 \) related RSs, and the reststrahlen band around the resonance where there are no RSs. (c) RSs over a larger frequency range, showing also the static mode with \( \omega_n = 0 \). (d) Relative error of RS frequencies calculated by the RSE.

C. Pole shift: Diagonal dRSE

To treat a shift in the pole positions of the permittivity, the idRSE requires the shifted pRSs to be included in the basis, along with the RSs of the unshifted poles. This can add a significant numerical effort, as the matrix diagonalization required to solve Eq. (9) has the computational complexity \( O(N^3) \). To avoid this, one can use a diagonal version of the dRSE, only retaining the diagonal elements in Eq. (9). We call this method ddRSE. It is linked to the first-order perturbation theory [18,35], which is also neglecting all off-diagonal elements. Accordingly, we expect this approach to be suited to treat small shifts in resonance frequency of the permittivity, such as the change of optical phonon frequencies with pressure or temperature. Neglecting the off-diagonal elements of Eq. (9) yields

\[
\omega = \omega_n \frac{1 - U_{nn}}{1 + V_{nn}}.
\]

(25)
The parameters of the GaAs microsphere and of the RSs shown are the same as in Fig. 3, with the pole of the permittivity shifting by 1% of its real part from $\tilde{\Omega}_1$ (black star) to $\Omega_1$ (blue star). Dispersive basis RSs (circles), exact RSs (squares), dRSE (crosses), and dRSE not including the infinitesimal-dispersive pRSs at $\tilde{\Omega}_1$; in the basis (diamonds), with $N = 315$ corresponding to $\Omega_1 \approx 0.4$ eV. (b) Relative error for the RSs calculated using the dRSE.

Considering only the resonance shift of one pole $\Omega_1$, without changing conductivities or $\varepsilon_{\infty}$, all $V_{nm}$ vanish, and

$$U_{nm} = \sum_{j=\pm 1} \left[ \frac{i}{\omega_n - \tilde{\Omega}_j} - \frac{i}{\omega_n - \Omega_j} \right] \int E_n(r) \cdot \hat{\sigma}_j(r) E_m(r) d^3r,$$

(26)

where $\tilde{\Omega}_1$ is the perturbed pole position.

To evaluate the suitability of this approach, we use the optical phonon-induced dispersion in GaAs discussed in Sec. IV B. We consider a shift of the phonon frequency by $1\%$, which is applied to the real part of $\Omega_1$, in order to obtain $\tilde{\Omega}_1$. We note that the GaAs phonon frequency under isotropic pressure shifts by 0.49 meV/GPa [36], so that a 1% change would correspond to exerting about 0.7 GPa pressure.

The results RSs close to $\Omega_1$ are shown in Fig. 4(a). The exact perturbed RSs converge towards the shifted pole $\tilde{\Omega}_1$, as expected. The diagonal (first order) approach is reproducing the exact RSs well when they are sufficiently far from the pole. Close to the pole instead, large errors are encountered. Notably, using all off-diagonal elements defined by Eq. (26) and solving the eigenvalue problem Eq. (9) does not significantly reduce the errors. This indicates that the effect of the basis incompleteness, missing the pRSs of $\tilde{\Omega}_1$, is dominating the remaining error.

The relative error of the dRSE results compared to the exact solutions is shown in Fig. 4(b). We find that the error for the WG modes is decreasing with increasing separation from $\Omega_1$. The leaky modes have a small error since they have little spatial overlap with the sphere, and are spectrally broad, making them insensitive to a shift of $\Omega_1$. We can thus conclude that the diagonal approximation, equivalent to a first-order perturbation theory, is a valid approach for small pole shifts, and separations of the RSs of interest from the poles much larger than these shifts.

The physical properties in scattering and absorption are expected to be dominated by the RSs away from the poles of the permittivity, as close to the poles the permittivity is very high, screening the outside field. Furthermore, for the response at real frequencies, the specific position of the RSs close to the poles is having a minor influence, since the poles typically have a significant imaginary part, and are therefore separated from the real axis.

D. Dispersive RSE: Absorption to gain

Another interesting feature one can address using the dRSE is the transition from absorption to gain, leading to lasing. In terms of the RSs, lasing occurs once the imaginary part of the RS frequency becomes positive, indicating an exponential increase of the field with time. This increase of the field comes with a corresponding increase of stimulated transitions, which in realistic systems results in gain saturation, reducing the imaginary part of the RS frequency to zero in a steady-state regime of lasing [29,37].

In the classical model of Lorentz oscillators, poles representing absorption have positive imaginary conductivity [23]. Continuously changing this conductivity to negative imaginary values transforms the absorption to gain. When representing the absorption due to interband transitions in a solid, a prefactor $1 - f_e - f_h$ is applied to the conductivity, where $f_e$ and $f_h$ are the occupation functions of, respectively, the related electron state in the conduction band and hole state in the valence band, which in thermal equilibrium are given by the Fermi distribution function [29]. Creating the population inversion changes the sign of the conductivity by changing the occupation factor $1 - f_e - f_h$ from 1 for no occupation ($f_e = f_h = 0$) to $-1$ for full occupation ($f_e = f_h = 1$).

As an example, we use GaAs, which is a direct band-gap semiconductor widely used in semiconductor lasers. To model the transition from absorption to gain, we concentrate on the energy region close to the band gap, where a population inversion is created by optical or electrical pumping, and fit the permittivity data [38] in the range 1.3 to 1.65 eV with four pairs of Lorentz poles having purely imaginary conductivities. The measured absorption index $n^\prime$ and the resulting fit of the Lorentz model are shown in Fig. 5(a). We see that the fit is not as good as for the optical-phonon range shown in Fig. 3(a), and allowing the conductivities to have nonzero real parts could lead to a better agreement, as in the case of gold. However, we intentionally fit the permittivity with a model of classical Lorentz oscillators, in order to enable a simulation of the population inversion. The fit function
contains three poles close to the band gap having a small linewidth to describe the sharp increase of the absorption at the band edge, and the fourth pole at higher energy outside the fit range, providing a background. When changing the sign of the first-pole conductivity \( \sigma_1 \), in order to simulate the change from no to full occupation, the resulting absorption index shows a negative region around 1.49 eV, indicating the region where the material exhibits gain.

We first consider a rather large microsphere of radius \( R = 10 \mu m \), and choose \( l = 10 \). The wavelength in the medium resonant to the gain is about \( \lambda = 0.24 \mu m \), much smaller than the radius, and thus the RSs close to the gain pole are of FP type for the chosen \( l \). To provide a round-trip amplification, the losses due to the transmission across the sphere surface must be compensated by the gain through the sphere. The amplitude reflection coefficient for normal incidence is \( r = (n'_r - 1)/(n'_r + 1) \approx 0.55 \), using the refractive index of \( n'_r = 3.5 \). The amplification of the wave amplitude propagating one
diameter is \( g = \exp(-2Rn'_r/\omega/c) \). The condition for round-trip amplification is then \( gr > 1 \), which requires \( n'_r < -0.004 \). We see in Fig. 5(a) that this condition is fulfilled around 1.49 eV.

The RSs under gain, calculated using the dRSE with the dispersive basis of RSs in the absorption regime (using a basis size \( N = 676 \) corresponding to \( \Omega_2 = 26.78 \) eV), are shown in Fig. 5(b) over the same energy range as used in Fig. 5(a). Away from the poles, the basis shows nearly equidistant FP modes, which have an imaginary part following the losses due to the absorption index \( n''_r \). In the absorption regime, close to the poles, the RSs converge from the lower frequency side towards the poles, which are at the transversal optical resonance frequencies \( \omega_\tau \), and show gaps at higher frequencies up to the longitudinal optical frequency \( \omega_L \), where \( n_r = 0 \), similar to the case of the optical phonons in Fig. 3(b).

Note that in the present case, the set of RSs approaching the longitudinal optical frequencies is rather large, on the order of 100. This is due to the large radial quantum number of the basis FP modes in the region of the pole frequency. This number can be estimated as \( 4R/\lambda \approx 166 \), using \( \lambda = 0.24 \mu m \).

After switching to gain, some RSs in the gain spectral region show a positive imaginary part, as expected from the above discussion, and are thus lasing RSs. Furthermore, a curious arrangement of RSs is observed close to the gain pole, which occurs due to the inverted influence of the pole—the refractive index now increases approaching the pole from higher frequencies, so that the infinite series of RSs now converges towards the pole from the higher frequency side. These RSs, however, form a loop, moving clockwise over the pole, starting at a value around the corresponding longitudinal optical frequency \( \omega_L \), which is now on the lower frequency side of the pole, as shown in the inset of Fig. 5(b). The original, unperturbed series of FP modes, present without the pole, is instead repelled from the pole. This is qualitatively different from the case of absorption, where the RS series is attracted to the pole, splitting the RSs into an infinite set of all radial quantum numbers below the pole frequency \( \omega_\tau = \Omega_2 \), and a second infinite set of all radial quantum numbers above \( \omega_L \).

To investigate the regime of WG mode lasing at \( l = 10 \), we consider in Fig. 6 a much smaller sphere of radius \( R = 0.94 \mu m \). In this case, the gain region lines up with a WG mode in the basis (WG3), moving it to a positive imaginary energy. The evolution of the imaginary part of WG3 with the change from absorption to gain is shown in the inset. We observe that with the build-up of the gain, given by the decreasing imaginary part of the conductivity, the imaginary part of the WG3 energy moves continuously towards positive values, eventually reaching the lasing regime. The other RSs only change slightly since they are not close to \( \Omega_1 \), so that WG3 remains the only lasing mode of the system.

V. CONCLUSIONS

We have shown a set of applications of the dispersive RSE, a method developed for accurate and efficient calculation of the resonant states of optical systems exhibiting frequency dispersion. We have also presented the infinitesimal-dispersive RSE, which uses the resonant states of a nondispersive system as a basis for treating an optical system with dispersion, adding new poles in the frequency dispersion in the limit of
nondispersive series, splitting it into two infinite series, above and below the pole. Adding a gain pole instead repels the nondispersive series, and creates an additional series forming a clockwise loop in complex frequency, starting from the longitudinal frequency left of the pole and converging towards the pole from the right.

Finally, we introduce a diagonal dispersive RSE, a first-order perturbation method for treating small changes in the pole position, which is suited for the resonant states sufficiently separated from the pole of the dispersion and which has a much lower numerical complexity than the full dispersive RSE.

We believe that the dispersive RSE, once implemented in an accessible software package, including arbitrary 3D open systems [18], and scattering and absorption with external excitation [32,39], will be a powerful and widely used tool for treating electromagnetic problems.

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APPENDIX A: DRUDE-LORENTZ MODEL OF THE PERMITTIVITY AND FIT FOR Au AND GaAs

Implementing causality, the dispersive permittivity obeys the relation $\hat{\epsilon}^\ast (\mathbf{r}, \omega) = \hat{\epsilon} (\mathbf{r}, -\omega^\ast)$. This implies that poles appear in pairs in the permittivity Eq. (2). In particular, each pole at $\Omega_j$ has a partner at $\Omega_{-j} = -\Omega_j^\ast$ with $\hat{\sigma}_{-j} = \hat{\sigma}_j^\ast$. A pole with $\text{Re} \Omega_j = 0$, such as that of the Drude model, can be seen as a pair of poles merged to produce a single pole on the imaginary $\omega$ axis with a purely real conductivity. Furthermore, Lorentz reciprocity requires that all tensors in Eq. (2) are symmetric. For isotropic materials, such as gold and GaAs, these tensors can be replaced by scalars.

In the calculations presented in this paper, we use for GaAs a Lorentz model of the permittivity consisting of $L$ pairs of poles,

$$\varepsilon(\omega) = \varepsilon_{\infty} + \sum_{j=1}^{L} \left( \frac{i\sigma_j}{\omega - \Omega_j} + \frac{i\sigma_j^\ast}{\omega + \Omega_j^\ast} \right),$$

and for gold a Drude-Lorentz model,

$$\varepsilon(\omega) = \varepsilon_{\infty} - \frac{\gamma \sigma}{\omega(\omega + i\gamma)} + \sum_{j=1}^{L} \left( \frac{i\sigma_j}{\omega - \Omega_j} + \frac{i\sigma_j^\ast}{\omega + \Omega_j^\ast} \right),$$

in which the Drude term of the dispersion has been added, consisting of Ohm’s law pole at $\omega = 0$ and a Drude pole at $\omega = \Omega_0 = -i\gamma$:

$$\frac{i\sigma}{\omega} - \frac{i\sigma}{\omega + i\gamma} = -\frac{\gamma \sigma}{\omega(\omega + i\gamma)}.$$  

Here $\varepsilon_{\infty}$ is the permittivity at high frequencies and $\sigma$ is the real DC conductivity. The generalized conductivities $\sigma_j = \sigma_j^\prime + i\sigma_j^\prime\prime$ at the Lorentz poles $\Omega_j = \Omega_j^\prime + i\Omega_j^\prime\prime$ are complex.

Table I contains the parameters of the fits shown in Figs. 1, 3, and 5, using the Drude-Lorentz and the Lorentz models with $L = 3, 1, \text{and } 4$, respectively. The resulting root-mean
TABLE I. Optimized model parameters for Au and GaAs phonon (*) and band-edge (**) ranges, using the fit function with optimization energy ranges corresponding to the data shown in Figs. 1, 3, 5, and 6. The number of data values \( N_d \) and the resulting error \( S \) are also given.

| Material | Au | GaAs* | GaAs** |
|----------|----|-------|--------|
| \( \epsilon_\infty \) | 0.5 | 11.0 | 8.6013 |
| \( \gamma \) (eV) | 0.065748 | – | – |
| \( \sigma \) (eV) | 1133.0 | – | – |
| \( \Omega_1 \) (eV) | 2.5936 | 0.033314 | 1.497 |
| \( \Omega_2 \) (eV) | -0.41875 | 1.4904 \( \times 10^{-4} \) | -0.03665 |
| \( \sigma_1 \) (eV) | 1.4029 | 0.0 | 0.0 |
| \( \sigma_2 \) (eV) | 0.76857 | 0.033262 | 0.01224 |
| \( \Omega_3 \) (eV) | 3.8192 | – | 1.5612 |
| \( \Omega_4 \) (eV) | -1.3246 | – | -0.05643 |
| \( \sigma_3 \) (eV) | 0.41939 | – | 0.0 |
| \( \sigma_4 \) (eV) | 4.5468 | – | 0.02432 |
| \( \Omega_5 \) (eV) | 9.8989 | – | 1.6463 |
| \( \Omega_6 \) (eV) | -4.2933 | – | -0.0457 |
| \( \sigma_5 \) (eV) | 0.012244 | – | 0.0 |
| \( \sigma_6 \) (eV) | 14.817 | – | 0.02404 |
| \( \Omega_7 \) (eV) | – | – | 2.2853 |
| \( \Omega_8 \) (eV) | – | – | -0.00778 |
| \( \sigma_7 \) (eV) | – | – | 0.0 |
| \( \sigma_8 \) (eV) | – | – | 2.9302 |
| \( h_{01} \) (eV) | 0.64 | 0.031 | 1.3 |
| \( h_{02} \) (eV) | 6.6 | 0.0372 | 1.65 |
| \( N_d \) | 49 | 47 | 8 |
| S | 1.4795 | 0.0372 | 0.0055 |

square error \( S \) of the fit is also shown. Its definition, with and without experimental errors taken into account in the optimization, is provided in Ref. [23]. This difference in the definition leads, in particular, to a much higher value of \( S \) for gold, in which case the fit takes the measurement errors [30] into account.

**APPENDIX B: TM MODES OF A SPHERE, THEIR NORMALIZATION AND MATRIX ELEMENTS**

The electric field of a TM mode of a sphere of radius \( R \) in vacuum has the form (in spherical coordinates)

\[
E(r) = \frac{A_{nl}}{n_r k_n r} \begin{pmatrix} l(l+1) \psi_{nl}(r) Y_{lm}(\theta, \varphi) \\
\frac{\partial}{\partial r} \psi_{nl}(r) \frac{\partial}{\partial \theta} Y_{lm}(\theta, \varphi) \\
\frac{1}{\sin \theta} \frac{\partial}{\partial \varphi} Y_{lm}(\theta, \varphi) \end{pmatrix},
\]

where \( Y_{lm}(\theta, \varphi) \) are real-valued spherical harmonics [18], and the radial wave functions within the sphere \( r \leq R \) are given by

\[
\psi_{nl}(r) = \frac{j_l(n_r k_n r)}{j_l(n_r k_n R)}.
\]

Here \( k_n = \omega_n/c \) is the RS wave number and \( n_r \) is the refractive index, which is given by

\[
n_r^2 = \varepsilon(\omega_n),
\]

with the permittivity \( \varepsilon(\omega) \). The normalization constant \( A_{nl} \) is found from the general RS normalization condition with dispersion [Eq. (8)] and has the form [7]

\[
\frac{1}{A_{nl}} = l(l+1)R^2 (n_r^2 - 1)n_r^2 D_{nl},
\]

with

\[
D_{nl} = \frac{1}{n_r^2} \left[ \frac{j_{l-1}(x)}{j_l(x)} - \frac{l}{x} - \frac{l(l+1)}{x^2} + \eta_n C_{nl} \right]
\]

and

\[
(n_r^2 - 1)C_{nl} = -2l \frac{j_{l-1}(x)}{x^2} + \frac{j_l(x)}{j_{l-2}(x)} - \frac{j_{l-2}(x)}{j_l(x)},
\]

where \( j_l(x) \) are the spherical Bessel functions, \( x = n_r \omega_n R/c \),

\[
\eta_n = \frac{\omega}{2k(\omega)} \left. \frac{\delta \varepsilon(\omega)}{\delta \omega} \right|_{\omega = \omega_n},
\]

and \( n_r \) is given by Eq. (B3).

The RS eigenfrequencies \( \omega_n \) are solutions of the secular equation for the TM polarization

\[
\frac{1}{n_r} \frac{j_{l-1}(n_r z)}{j_l(n_r z)} = \frac{h_{l-1}(z)}{h_l(z)} - \frac{l}{z} \left( 1 - \frac{1}{n_r^2} \right),
\]

in which \( h_l(z) \equiv h_l^{(1)}(z) \) is the spherical Hankel function of the first kind, with \( z = \omega_n R/c \), and \( n_r(\omega_n) \) is defined by Eq. (B3).

For small values of \( R \), corresponding to the electrostatic limit, the spherical Bessel and Hankel functions can be expanded into power series of their arguments and keeping only the leading terms, which is valid for \( |\omega| R/c \ll 1 \) and \( |n_r(\omega)\omega R/c | \ll 1 \), one can obtain from Eq. (B8) a well-known equation

\[
\varepsilon(\omega) = -\frac{l+1}{l},
\]

determining for each \( l \) the eigenfrequency \( \omega \) of the surface plasmon in the low-frequency (electrostatic) limit.

For the pRRs, due to a pole of the dispersion at \( \Omega_j \), all the eigenfrequencies are the same, \( \omega_n = \Omega_j \), but the eigenvalues of the refractive index \( n_r(\omega_n) \) are different. They are again determined by the secular equation (B8) which is now solved for \( n_r \), while keeping \( z = \Omega_j R/c \) fixed. These eigenvalues \( n_r \) are linked to \( q_n \), as defined by Eq. (14),

\[
n_r^2(\omega_n) = \varepsilon_\infty + \sum_{j \neq j} \frac{i \sigma_j}{\Omega_j - \Omega_j} + \frac{1}{q_n},
\]

and to the values \( \eta_n \) in the normalization constants Eq. (B7):

\[
\eta_n = \frac{1}{\xi} \frac{\Omega_j}{2n_r^2 q_n^2} = -\frac{1}{\alpha^2} \frac{1}{2n_r^2 q_n},
\]

in accordance with Eqs. (15) and (17), using \( \xi = \alpha^2 \Omega_j/q_n \). Clearly \( \xi \to 0 \) makes the normalization constants \( A_{nl} \) vanishing. Redefining these constants as

\[
A_{nl} = \alpha_a \tilde{A}_{nl},
\]

with

\[
\tilde{A}_{nl} = \frac{1}{n_r^2} \left[ \frac{j_{l-1}(x)}{j_l(x)} - \frac{l}{x} - \frac{l(l+1)}{x^2} + \eta_n C_{nl} \right]
\]

and

\[
(n_r^2 - 1)C_{nl} = -2l \frac{j_{l-1}(x)}{x^2} + \frac{j_l(x)}{j_{l-2}(x)} - \frac{j_{l-2}(x)}{j_l(x)},
\]
in accordance with Eq. (18), we find the new (finite) normalization constants of the pRSs:

$$\frac{1}{A_{nl}} = -l(l+1)R^3(n_z^2 - 1) \frac{1}{2q_n} C_{nl}.$$  \hspace{1cm} (B13)

For any homogeneous perturbation of the sphere, all the matrix elements contributing to Eq. (9) or Eq. (20) are given by the integrals

$$W_{nm} = \int_{V_R} \mathbf{\hat{E}}_n(r) \cdot \mathbf{\hat{E}}_m(r) \, dr,$$  \hspace{1cm} (B14)

where $V_R$ is the volume of the sphere. These integrals have the following analytic form:

$$W_{nm} = \tilde{A}_{nl} \tilde{A}_{ml}(l+1)R^3 F_l(x, y),$$  \hspace{1cm} (B15)

where $x = n_r(\omega_0) \alpha_0 R/c$ and $y = n_r(\omega_0) \alpha_0 R/c$, and both RSs $n$ and $m$ belong to the same orbital quantum number $l$. The function $F_l(x, y)$ is defined as

$$F_l(x, y) = \frac{1}{x^2 - y^2}\left[\frac{xj_l(y)}{j_l(x)} - \frac{yj_l(x)}{j_l(y)}\right] - \frac{l}{xy},$$  \hspace{1cm} (B16)

which for coinciding arguments becomes

$$F_l(x, x) = \frac{1}{2}\left[\frac{j_l'(x)}{j_l(x)} - \frac{j_l'(x)}{j_l(x)}\right] - \frac{l}{x^2}.$$  \hspace{1cm} (B17)

Note that using the normalization coefficients Eq. (B13) for the pRSs, the diagonal elements Eq. (B15) become $W_{nn} = -q_n$, in accordance with their normalization condition

$$1 = -\frac{1}{q_n} \int_{V_R} |\mathbf{\hat{E}}_n(r)|^2 \, dr,$$  \hspace{1cm} (B18)

see Eq. (16).

**APPENDIX C: CALCULATING THE BASIS RSs**

To generate a basis of unperturbed RSs, the secular equation (B8) is solved numerically using the Newton-Raphson algorithm. This relies on a suited choice of guess values, which for $|n_z, z| \lesssim 1$ are found on a two-dimensional equidistant grid in the complex $z$ plane. For large values of $n_z, z$

however, we use approximate recursive relations given below, which provide reliable guess values based on the eigenvalues found for the neighboring RSs. To derive these relations, it is convenient to modify Eq. (B8) to the following form:

$$\frac{1}{n_r} \frac{j'_l(n_r z)}{j_l(n_r z)} = \frac{\bar{h}'_l(z)}{\bar{h}_l(z)} + \frac{1}{z} \left(1 - \frac{1}{n_r^2}\right),$$  \hspace{1cm} (C1)

where $j'_l(x)$ and $\bar{h}'_l(x)$ are the derivatives of the spherical Bessel and Hankel functions with respect to their arguments. Noting that

$$j_l(x) \approx \frac{1}{x} \cos \left(x - \frac{l + 1}{2}\right),$$  \hspace{1cm} (C2)

at large $x$, we find

$$\frac{j'_l(x)}{j_l(x)} \approx -\tan \left(x - \frac{l + 1}{2}\right),$$  \hspace{1cm} (C3)

which provides an estimate for the left-hand side of Eq. (C1). Its right-hand side is in turn a slowly changing function. Therefore, the neighboring RSs are separated by $\Delta z \approx \pi$, which is the period of the tangent function. This separation can be further evaluated as

$$\Delta z = n_r(\Delta z + \Delta z) - n_r(\Delta z) \approx (n_r + n'_r) \Delta z,$$  \hspace{1cm} (C4)

where $n'_r = dn_r/dz$. Then we find approximate relations for the neighboring roots of Eq. (B8) in the following three cases:

(i) No dispersion of permittivity or RS eigenfrequencies are far from any poles of the dispersion, i.e., $n_r$ is nearly constant:

$$\Delta z \approx \frac{\pi}{n_r},$$  \hspace{1cm} (C5)

(ii) Strong dispersion, such as that for the RS near the pole of the dispersion:

$$\Delta z \approx \frac{\pi}{n_r + n'_r},$$  \hspace{1cm} (C6)

(iii) Pole RSs—which are required for constructing an infinitesimal-dispersive basis. In this case Eq. (B8) is solved for $n_r$ with $z$ fixed. For the neighboring pRSs, we find

$$\Delta n_r \approx \frac{\pi}{z}.$$  \hspace{1cm} (C7)

[1] G. Gamow, Z. Phys. **51**, 204 (1928).
[2] A. F. J. Siegert, Phys. Rev. **56**, 750 (1939).
[3] E. A. Muljarov, W. Langbein, and R. Zimmermann, Europhys. Lett. **92**, 50010 (2010).
[4] C. Sauvan, J. P. Hugonin, I. S. Maksymov, and P. Lalanne, Phys. Rev. Lett. **110**, 237401 (2013).
[5] Q. Bai, M. Perrin, C. Sauvan, J.-P. Hugonin, and P. Lalanne, Opt. Express **21**, 27371 (2013).
[6] M. Perrin, Opt. Express **24**, 27137 (2016).
[7] E. A. Muljarov and W. Langbein, Phys. Rev. B **94**, 235438 (2016).
[8] E. A. Muljarov and T. Weiss, Opt. Lett. **43**, 1978 (2018).
[9] E. A. Muljarov and W. Langbein, Phys. Rev. A **96**, 017801 (2017).
[10] P. Lalanne, W. Yan, K. Vynck, C. Sauvan, and J.-P. Hugonin, Laser Photon. Rev. **12**, 1700113 (2018).
[11] P. L. Kapur and R. Peierls, Proc. R. Soc. London Ser. A **166**, 277 (1938).
[12] D. J. Bergman, Phys. Rev. B **19**, 2359 (1979).
[13] H. E. Türeci, A. D. Stone, and B. Collier, Phys. Rev. A **74**, 043822 (2006).
[14] J. Mikkalou, M. Kauranen, and S. Suuriniemi, Phys. Rev. B **89**, 165429 (2014).
[15] P. Y. Chen, D. J. Bergman, and Y. Sivan, Phys. Rev. Appl. **11**, 044018 (2019).
[16] G. Arfken and H. Weber, Mathematical Methods for Physicists (Elsevier, New York, 2013).
[17] M. B. Doost, W. Langbein, and E. A. Muljarov, Phys. Rev. A **87**, 043827 (2013).
[18] M. B. Doost, W. Langbein, and E. A. Muljarov, Phys. Rev. A 90, 013834 (2014).
[19] W. Yan, R. Faggiani, and P. Lalanne, Phys. Rev. B 97, 205422 (2018).
[20] L. Zschiedrich, F. Binkowski, N. Nikolay, O. Benson, G. Kewes, and S. Burger, Phys. Rev. A 98, 043806 (2018).
[21] P. Lalanne, W. Yan, A. Gras, C. Sauvan, J.-P. Hugonin, M. Besbes, G. Demésy, M. D. Truong, B. Gralak, F. Zolla et al., J. Opt. Soc. Am. A 36, 686 (2019).
[22] E. A. Muljarov and W. Langbein, Phys. Rev. B 93, 075417 (2016).
[23] H. S. Sehmi, W. Langbein, and E. A. Muljarov, Phys. Rev. B 95, 115444 (2017).
[24] H. S. Sehmi, W. Langbein, and E. A. Muljarov, in 2017 Progress In Electromagnetics Research Symposium - Spring (PIERS) (IEEE, New York, 2017), pp. 994–1000.
[25] S. V. Lobanov, G. Zoriniants, W. Langbein, and E. A. Muljarov, Phys. Rev. A 95, 053848 (2017).
[26] T. Shahbazyan and M. Stockman, Plasmonics: Theory and Applications (Springer, Netherlands, 2014).
[27] K. Vahala, Optical Microcavities (World Scientific, Singapore, 2004).
[28] A. Kavokin, J. J. Baumberg, G. Malpuech, and F. P. Laussy, Microcavities (Oxford University Press, Oxford, 2007).
[29] W. W. Chow, S. W. Koch, and M. I. I. I. Sargent, Semiconductor-Laser Physics (Springer, Berlin, 2012).
[30] P. B. Johnson and R. W. Christy, Phys. Rev. B 6, 4370 (1972).
[31] C. F. Bohren and D. R. Huffman, Absorption and Scattering of Light by Small Particles (Wiley-VCH, New York, 1998).
[32] S. V. Lobanov, W. Langbein, and E. A. Muljarov, Phys. Rev. A 98, 033820 (2018).
[33] V. Myroshnychenko, J. Rodríguez-Fernández, I. Pastoriza-Santos, A. M. Funston, C. Novo, P. Mulvaney, L. M. Liz-Marzán, and F. J. G. de Abajo, Chem. Soc. Rev. 37, 1792 (2008).
[34] J. S. Blakemore, J. Appl. Phys. 53, 123 (1982).
[35] T. Weiss, M. Mesch, M. Schäferling, H. Giessen, W. Langbein, and E. A. Muljarov, Phys. Rev. Lett. 116, 237401 (2016).
[36] S. Adachi, GaAs and Related Materials: Bulk Semiconducting and Superlattice Properties (World Scientific, Singapore, 1994).
[37] H. E. Türeci, L. Ge, S. Rotter, and A. D. Stone, Science 320, 643 (2008).
[38] The data for the measured indices of light refraction and absorption is taken from http://sspectra.com/sopra.html.
[39] T. Weiss and E. A. Muljarov, Phys. Rev. B 98, 085433 (2018).